

APPROVED DRUG PRODUCTS

WITH

**THERAPEUTIC
EQUIVALENCE
EVALUATIONS**

45th EDITION

**THE PRODUCTS IN THIS LIST HAVE BEEN APPROVED UNDER
SECTION 505 OF THE FEDERAL FOOD, DRUG, AND COSMETIC ACT.**

**U.S. DEPARTMENT OF HEALTH AND HUMAN SERVICES
FOOD AND DRUG ADMINISTRATION
OFFICE OF MEDICAL PRODUCTS AND TOBACCO
CENTER FOR DRUG EVALUATION AND RESEARCH
OFFICE OF GENERIC DRUGS
OFFICE OF GENERIC DRUG POLICY**

2025

APPROVED DRUG PRODUCTS with **THERAPEUTIC EQUIVALENCE EVALUATIONS**

The products in this list have been approved under section 505 of the Federal Food, Drug, and Cosmetic Act. This volume is current through December 31, 2024.

45th EDITION



U.S. DEPARTMENT OF HEALTH AND HUMAN SERVICES
FOOD AND DRUG ADMINISTRATION
OFFICE OF MEDICAL PRODUCTS AND TOBACCO
CENTER FOR DRUG EVALUATION AND RESEARCH
OFFICE OF GENERIC DRUGS
OFFICE OF GENERIC DRUG POLICY

2025

**FOOD AND DRUG ADMINISTRATION
CENTER FOR DRUG EVALUATION AND RESEARCH**

**APPROVED DRUG PRODUCTS
With
Therapeutic Equivalence Evaluations**

CONTENTS

	<i>PAGE</i>
PREFACE TO FORTY FIFTH EDITION.....	iv
1.0 INTRODUCTION.....	vi
1.1 Content and Exclusion.....	vi
1.2 Therapeutic Equivalence-Related Terms.....	vii
1.3 Further Guidance on Bioequivalence.....	ix
1.4 RLD and Reference Standard.....	x
1.5 General Policies and Legal Status.....	xi
1.6 Practitioner/User Responsibilities.....	xi
1.7 Therapeutic Equivalence Evaluations Codes.....	xii
1.8 Description of Certain Special Situations.....	xx
1.9 Therapeutic Equivalence Code Change for a Category of Multisource Drug Products.....	xxiii
1.10 Change of the Therapeutic Equivalence Evaluation for a Single Product.....	xxiii
1.11 Discontinued Drug Product List.....	xxiv
1.12 Changes to the Orange Book.....	xxiv
1.13 Availability of the Edition.....	xxv
2.0 HOW TO USE THE DRUG PRODUCTS LISTS.....	2-1
2.1 Key Sections for Using the Drug Product Lists.....	2-1
2.2 Drug Product Illustration.....	2-3
2.3 Therapeutic Equivalence Evaluations Illustration.....	2-4
DRUG PRODUCT LISTS	
Prescription Drug Product List.....	3-1
OTC Drug Product List.....	4-1
Drug Products with Approval under Section 505 of the FD&C Act Administered by the Center for Biologics Evaluation and Research List.....	5-1
Discontinued Drug Product List.....	6-1
Orphan Products Designations and Approvals List.....	7-1
Drug Products Which Must Demonstrate in vivo Bioavailability Only if Product Fails to Achieve Adequate Dissolution.....	8-1
APPENDICES	
A. Product Name Index.....	A-1
B. Product Name Index Listed by Applicant.....	B-1
C. Uniform Terms.....	C-1
PATENT AND EXCLUSIVITY INFORMATION ADDENDUM.....	
A. Patent and Exclusivity Lists.....	ADA1
B. Patent and Exclusivity Terms.....	ADB1

**FOOD AND DRUG ADMINISTRATION
CENTER FOR DRUG EVALUATION AND RESEARCH**

**APPROVED DRUG PRODUCTS
With
Therapeutic Equivalence Evaluations**

PREFACE TO FORTY FIFTH EDITION

The publication, *Approved Drug Products With Therapeutic Equivalence Evaluations* (the List, commonly known as the Orange Book), identifies drug products approved on the basis of safety and effectiveness by the Food and Drug Administration (FDA) under the Federal Food, Drug, and Cosmetic Act (the FD&C Act). The main criterion for the inclusion of any product is that the product is the subject of an application with an approval that has not been withdrawn for safety or effectiveness reasons. Inclusion of products in the Orange Book is independent of any current regulatory action being taken administratively or judicially against a drug product. In addition, the Orange Book contains therapeutic equivalence evaluations for approved multisource prescription drug products. These evaluations have been prepared to serve as public information and advice to state health agencies, prescribers, and pharmacists to promote public education in the area of drug product selection and to foster containment of healthcare costs. Therapeutic equivalence evaluations in this publication are not official FDA actions affecting the legal status of products under the FD&C Act.

Background of the Publication. To contain drug costs, virtually every state has adopted laws and/or regulations that encourage the substitution of drug products. These state laws generally require either that substitution be limited to drugs on a specific list (the positive formulary approach) or that it be permitted for all drugs except those prohibited by a particular list (the negative formulary approach). Because of the number of requests in the late 1970s for FDA assistance in preparing both positive and negative formularies, it became apparent that FDA could not serve the needs of each state on an individual basis. The Agency also recognized that providing a single list based on common criteria would be preferable to evaluating drug products on the basis of differing definitions and criteria in various state laws. As a result, on May 31, 1978, the Commissioner of the Food and Drug Administration sent a letter to officials of each state announcing FDA's intent to provide a list of all prescription drug products that are approved by FDA for safety and effectiveness, along with therapeutic equivalence determinations for multisource prescription products.

The Orange Book was distributed as a proposal in January 1979. It included only currently marketed prescription drug products approved by FDA through new drug applications (NDAs) and abbreviated new drug applications (ANDAs) under the provisions of Section 505 of the FD&C Act and FDA regulations at that time.

The therapeutic equivalence evaluations in the Orange Book reflect FDA's application of specific criteria to the multisource prescription drug products listed in the Orange Book and approved under Section 505 of the FD&C Act. These evaluations are presented in the form of code letters that indicate the basis for the evaluation made. An explanation of the codes appears in the *Introduction*.

A complete discussion of the background and basis of FDA's therapeutic equivalence evaluation policy was published in the *Federal Register* on January 12, 1979 (44 FR 2932). The final rule, which includes FDA's responses to the public comments on the proposal, was published in the *Federal Register* on October 31, 1980 (45 FR 72582). The first publication of the Orange Book in October 1980, concurrent with finalization of the rule, incorporated appropriate revisions. Each subsequent edition has included new approvals and made appropriate changes in data.

On September 24, 1984, the President signed into law the Drug Price Competition and Patent Term Restoration Act of 1984 (Hatch-Waxman Amendments). The Hatch-Waxman Amendments amended the FD&C Act to establish, among other things, the 505(b)(2) and 505(j) approval pathways. The Hatch-Waxman Amendments require that FDA, among other things, make publicly available a list of approved drug products with monthly supplements. The Orange Book and its monthly Cumulative Supplements satisfy this requirement. The *Addendum* to this publication identifies drugs that have qualified under the FD&C Act for periods of exclusivity and provides patent information concerning the approved drug products in the Orange Book. The *Addendum* also provides additional information that may be helpful to those submitting an NDA under Section 505(b) of the FD&C Act or an ANDA under Section 505(j) of the FD&C Act to the Agency.

The Agency intends to use this publication to further its objective of obtaining input and comment on the publication itself and related Agency procedures. Therefore, if you have comments on how the publication can be improved, please send them to the Central Document Room, Attn: Director, Division of Orange Book Publication and Regulatory Assessment (DOBPR), Office of Generic Drug Policy, Center for Drug Evaluation and Research, Food and Drug Administration, 5901-B Ammendale Rd., Beltsville, MD 20705-1266. Comments received are publicly available to the extent allowable under the Freedom of Information Act and FDA regulations.

1.0 INTRODUCTION

1.1 Content and Exclusion

The Orange Book is composed of four parts: (1) approved prescription drug products with therapeutic equivalence evaluations (Prescription Drug Product List); (2) approved Over-the-Counter (OTC) drug products (OTC Drug Product List); (3) Drug Products with Approval under Section 505 of the FD&C Act administered by the Center for Biologics Evaluation and Research List; and (4) a cumulative list of approved products that have never been marketed, are for exportation, are for military use, have been discontinued from marketing and we have not determined that they were withdrawn from sale for safety or effectiveness reasons, or have had their approvals withdrawn for other than safety or effectiveness reasons subsequent to being discontinued from marketing (Discontinued Drug Product List).¹ This publication also includes Appendices of information from prescription and OTC drug products lists:

- (1) Appendix A, which is an alphabetical list of proprietary names, if one exists (brand name or trade name), and the name of the active ingredient.
- (2) Appendix B, which is an alphabetical list of applicant names which have been abbreviated for this publication.
- (3) Appendix C, which is an alphabetical list of uniform terms.

The *Addendum* contains patent and exclusivity information for the products contained in the Prescription Drug Product, OTC Drug Product, Discontinued Drug Product, and the Drug Products with Approval under Section 505 of the FD&C Act administered by the Center for Biologics Evaluation and Research Lists. The publication may include additional information that the Agency deems appropriate to disseminate.

Prior to the 6th Edition, the publication had excluded OTC drug products and drug products with approval under Section 505 of the FD&C Act administered by the Center for Biologics Evaluation and Research. The Hatch-Waxman Amendments required the Agency to begin publishing an up-to-date list of all marketed drug products, OTC as well as prescription, that have been approved for safety and effectiveness and for which NDAs are required.

Under the FD&C Act, some drug products are given tentative approvals. The Agency will not include drug products with tentative approvals in the Orange Book because a drug product that is granted tentative approval is not an approved drug product. Tentative approval lists by month are available on FDA's website Drugs@FDA. When the tentative approval becomes a final approval through a subsequent action letter to the applicant, the Agency will list the drug product and the date of approval in the appropriate approved drug product list. In addition, we note that Section 505(x) of the FD&C Act affects the date of approval for certain drug products subject to scheduling under the Controlled Substances Act. For these drug products subject to scheduling, the Agency will list the drug product on FDA's website Drugs@FDA upon approval under Section 505(c) of the FD&C Act, and will list the drug product in the Orange Book upon the date of approval as determined under Section 505(x).

¹ Generally, newly approved products are added to the Active Section of the Orange Book (i.e., the Prescription Drug Product List or the OTC Drug Product List), depending on the dispensing requirements (prescription or OTC) or approval authority, unless the Division of Orange Book Publication and Regulatory Assessment is otherwise notified before publication. See Section 1.12.

The Orange Book identifies the application holder of a drug product and does not identify distributors or repackagers.

1.2 Therapeutic Equivalence-Related Terms

Pharmaceutical Equivalents. Pharmaceutical equivalents are drug products in identical dosage forms and route(s) of administration that contain identical amounts of the identical active drug ingredient, i.e., the same salt or ester of the same therapeutic moiety, or, in the case of modified-release dosage forms that require a reservoir or overage or such forms as prefilled syringes where the residual volume may vary, that deliver identical amounts of the active drug ingredient over the identical dosing period; do not necessarily contain the same inactive ingredients; and meet the identical compendial or other applicable standard of identity, strength, quality, and purity, including potency and, where applicable, content uniformity, disintegration times, and/or dissolution rates.² They may differ in characteristics such as shape, scoring configuration, release mechanisms, packaging, excipients (including colors, flavors, preservatives), expiration date/time, and, within certain limits, labeling.

Pharmaceutical Alternatives. Pharmaceutical alternatives are drug products that contain the identical therapeutic moiety, or its precursor, but not necessarily in the same amount or dosage form, or the same salt or ester (e.g., tetracycline hydrochloride, 250 mg capsules vs. tetracycline phosphate complex, 250 mg capsules; quinidine sulfate, 200 mg tablets vs. quinidine sulfate, 200 mg capsules).³ Each such drug product individually meets either the identical or its own respective compendial or other applicable standard of identity, strength, quality, and purity, including potency and, where applicable, content uniformity, disintegration times, and/or dissolution rates.⁴ Different dosage forms and strengths within a product line by a single manufacturer are pharmaceutical alternatives, as are extended-release products when compared with immediate-release or standard-release formulations of the same active ingredient.

Therapeutic Equivalents. Approved drug products are considered to be therapeutic equivalents if they are pharmaceutical equivalents for which bioequivalence has been demonstrated, and they can be expected to have the same clinical effect and safety profile when administered to patients under the conditions specified in the labeling.⁵

FDA classifies as therapeutically equivalent those drug products that meet the following general criteria: (1) they are approved as safe and effective; (2) they are pharmaceutical equivalents in that they (a) contain identical amounts of the identical active drug ingredient in the identical dosage form and route of administration, and (b) meet compendial or other applicable standards of strength, quality, purity, and identity; (3) they are bioequivalent in that (a) they do not present a known or potential bioequivalence problem, and they may meet an acceptable *in vitro* standard, or (b) if they do present such a known or potential problem, they are shown to meet an appropriate bioequivalence standard (4) they are adequately labeled; and (5) they are manufactured in compliance with Current Good Manufacturing Practice regulations. The concept of therapeutic equivalence applies only to

² 21 CFR 314.3(b).

³ See 21 CFR 314.3(b).

⁴ 21 CFR 314.3(b).

⁵ 21 CFR 314.3(b).

drug products containing the identical active ingredient(s) and does not encompass a comparison of different therapeutic agents used for the same condition (e.g., meperidine hydrochloride vs. morphine sulfate for the treatment of pain). Any drug product in the Orange Book repackaged and/or distributed by other than the applicant is considered to be therapeutically equivalent to the applicant's drug product even if the applicant's drug product is single source or coded as non-equivalent (e.g., **BN**). Distributors or repackagers of an applicant's drug product are not identified in the Orange Book.

FDA considers drug products to be therapeutically equivalent if they meet the criteria outlined above, even though they may differ in certain other characteristics such as shape, scoring configuration, release mechanisms, packaging, excipients (including colors, flavors, preservatives), expiration date/time, certain aspects of labeling (e.g., the presence of specific pharmacokinetic information), and storage conditions. When such differences are important in the care of a particular patient, it may be appropriate for the prescribing physician to require that a specific product be dispensed as a medical necessity. With this limitation, however, FDA believes that products classified as therapeutically equivalent can be substituted with the full expectation that the substituted product can be expected to have the same clinical effect and safety profile as the prescribed product when administered to patients under the conditions specified in the labeling.

Strength. Strength refers to the amount of drug substance contained in, delivered, or deliverable from a drug product, which includes: (1)(a) the total quantity of drug substance in mass or units of activity in a dosage unit or container closure (e.g., weight/unit dose, weight/volume or weight/weight in a container closure, or units/volume or units/weight in a container closure); and/or, as applicable, (b) the concentration of the drug substance in mass or units of activity per unit volume or mass (e.g., weight/weight, weight/volume, or units/volume); or (2) such other criteria the Agency establishes for determining the amount of drug substance contained in, delivered, or deliverable from a drug product if the weights and measures described in clause (1)(a) do not apply (e.g., certain drug-device combination products for which the amount of drug substance is emitted per use or unit time).⁶ Note that if the criteria the Agency establishes for determining and expressing the amount of drug substance in a product evolves over time, the Agency generally does not intend to revise the expressions of strength for drug products already included in the Orange Book, but rather intends to apply the criteria prospectively to drug products added to the Orange Book.

Although the strength of drug products in the Orange Book is generally expressed in terms of the amount of drug substance (active ingredient) in the drug product, it is sometimes expressed in terms of the amount of the active moiety.⁷ For example, certain drug products included in the Orange Book include a designation of "EQ" next to their expression of strength. This "EQ" designation generally is used in connection with salt drug products to indicate that the strength of such drug product is being expressed in terms of the equivalent strength of the active moiety (e.g., "EQ 200 MG BASE"), rather than in terms of the strength of the active ingredient.

Bioavailability. Bioavailability is the rate and extent to which the active ingredient or active moiety is absorbed from a drug product and

⁶ 21 CFR 314.3(b).

⁷ Active moiety is the molecule or ion, excluding those appended portions of the molecule that cause the drug to be an ester, salt (including a salt with hydrogen or coordination bonds), or other noncovalent bonds (such as a complex, chelate, or clathrate) of the molecule, responsible for the physiological or pharmacological action of the drug substance. 21 CFR 314.3(b).

becomes available at the site of drug action. For drug products that are not intended to be absorbed into the bloodstream, bioavailability may be assessed by scientifically valid measurements intended to reflect the rate and extent to which the active ingredient or active moiety becomes available at the site of drug action.⁸

Bioequivalence. Bioequivalence is the absence of a significant difference in the rate and extent to which the active ingredient or active moiety in pharmaceutical equivalents or pharmaceutical alternatives becomes available at the site of drug action when administered at the same molar dose under similar conditions in an appropriately designed study.⁹ Section 505(j)(8)(B) of the FD&C Act describes certain conditions under which a test drug and reference listed drug (RLD)(see Section 1.4) shall be considered bioequivalent:

- (i) the rate and extent of absorption of the [test] drug do not show a significant difference from the rate and extent of absorption of the [reference] listed drug when administered at the same molar dose of the therapeutic ingredient under similar experimental conditions in either a single dose or multiple doses; or
- (ii) the extent of absorption of the [test] drug does not show a significant difference from the extent of absorption of the [reference] listed drug when administered at the same molar dose of the therapeutic ingredient under similar experimental conditions in either a single dose or multiple doses and the difference from the [reference] listed drug in the rate of absorption of the drug is intentional, is reflected in its proposed labeling, is not essential to the attainment of effective body drug concentrations on chronic use, and is considered medically insignificant for the drug.

Where these above methods are not applicable (e.g., for drug products that are not intended to be absorbed into the bloodstream), other scientifically valid *in vivo* or *in vitro* test methods to demonstrate bioequivalence may be appropriate.

For example, bioequivalence may sometimes be demonstrated using an *in vitro* bioequivalence standard, especially when such an *in vitro* test has been correlated with human *in vivo* bioavailability data. In other situations, bioequivalence may sometimes be demonstrated through comparative clinical trials or pharmacodynamic studies.¹⁰

1.3 Further Guidance on Bioequivalence

FDA's regulations and guidance documents provide additional information regarding bioequivalence and bioavailability, including methodologies and statistical criteria used to establish the bioequivalence of drug products.¹¹

⁸ 21 CFR 314.3(b).

⁹ 21 CFR 314.3(b).

¹⁰ 21 CFR 320.24.

¹¹ We note that prior to the 36th edition of the Orange Book, the Preface to the Orange Book included a section entitled "Statistical Criteria for Bioequivalence." Please see FDA's regulations and guidance documents for additional information regarding bioequivalence and bioavailability. See generally 21 CFR part 320. See FDA Drugs guidance Web page at <https://www.fda.gov/drugs/guidance-compliance-regulatory-information/guidances-drugs> and FDA Drugs guidance (Product-Specific Guidances for Generic Drug Development) Web page at <https://www.fda.gov/drugs/guidances-drugs/product-specific-guidances-generic-drug-development>.

1.4 RLD and Reference Standard

An RLD is the listed drug¹² identified by FDA as the drug product upon which an applicant relies in seeking approval of its ANDA.¹³ FDA's general practice is to designate as RLDs drug products that have been approved for safety and effectiveness under Section 505(c) of the FD&C Act. For an ANDA based on an approved suitability petition (a petitioned ANDA), the RLD generally is the listed drug referenced in the approved suitability petition.¹⁴

A reference standard is the drug product selected by FDA that an applicant seeking approval of an ANDA must use in conducting an *in vivo* bioequivalence study required for approval.¹⁵ FDA generally selects a single reference standard that ANDA applicants must use in *in vivo* bioequivalence testing. Ordinarily, FDA will select the RLD as the reference standard. However, in some instances, the RLD and the reference standard may be different. For example, where the RLD has been withdrawn from sale for reasons other than safety or effectiveness, FDA may select an ANDA that is therapeutically equivalent to this RLD as the reference standard.

FDA identifies RLDs in the Prescription Drug Product, OTC Drug Product, and Discontinued Drug Product Lists. Listed drugs identified as RLDs represent drug products upon which an applicant can rely in seeking approval of an ANDA. FDA intends to update periodically the RLDs identified in the Prescription Drug Product, OTC Drug Product, and Discontinued Drug Product Lists, as appropriate.

If FDA has not designated an RLD for a drug product the applicant intends to duplicate, the potential applicant may submit a controlled correspondence to the Office of Generic Drugs to ask FDA to designate an RLD for that drug product. Section 1.7, *Therapeutic Equivalence Evaluations Codes (products meeting necessary bioequivalence requirements)*, explains the character coding system (e.g., **AB**, **AB1**, **AB2**, **AB3**...) for multisource prescription drug products listed under the same heading with two or more RLDs.

FDA also identifies reference standards in the Prescription Drug Product and OTC Drug Product Lists. Listed drugs identified as reference standards represent FDA's best judgment at this time as to the appropriate comparator for purposes of conducting any *in vivo* bioequivalence studies required for approval.

A potential applicant should consult Agency guidance related to referencing approved drug products in ANDA submissions for information on submitting a request for selection of a reference standard. FDA may, on its own initiative, select a new reference standard when doing so will help to ensure that applications for generic drugs may be submitted and evaluated, e.g., in the event that the listed drug currently selected as the reference standard has been withdrawn from sale.

¹² A "listed drug" is a new drug product that has been approved under Section 505(c) of the FD&C Act for safety and effectiveness or under Section 505(j) of the FD&C Act, which has not been withdrawn or suspended under Section 505(e)(1) through (5) or Section 505(j)(6) of the FD&C Act, and which has not been withdrawn from sale for what FDA has determined are reasons of safety or effectiveness. Listed drug status is evidenced by the drug product's identification in the current edition of FDA's "Approved Drug Products With Therapeutic Equivalence Evaluations" (Orange Book or the list) as an approved drug. A drug product is deemed to be a listed drug on the date of approval for the NDA or ANDA for that drug product (21 CFR 314.3(b)).

¹³ 21 CFR 314.3(b).

¹⁴ 21 CFR 314.94(a)(3)(i).

¹⁵ 21 CFR 314.3(b).

If an applicant has a question related to the appropriate reference standard, it is recommended that an applicant submit a controlled correspondence to the Office of Generic Drugs.

1.5 General Policies and Legal Status

The Orange Book contains public information and advice. It does not mandate the drug products that are purchased, prescribed, dispensed, or substituted for one another, nor does it, conversely, mandate the products that should be avoided. To the extent that the Orange Book sets forth FDA's evaluations of the therapeutic equivalence of drug products that have been approved, it contains FDA's advice to the public, to practitioners, and to the states regarding drug product selection. These evaluations do not constitute determinations that any product is in violation of the FD&C Act or that any product is preferable to any other. Therapeutic equivalence evaluations are a scientific judgment based upon evidence, while generic substitution may involve social and economic policy administered by the states, e.g., reducing the cost of drugs to consumers. To the extent that the Orange Book identifies drug products approved under Section 505 of the FD&C Act, it sets forth information that the Agency is required to publish and that the public is entitled to under the Freedom of Information Act. Exclusion of a drug product from the Orange Book does not necessarily mean that the drug product is in violation of Section 505 of the FD&C Act, that such a product is not safe or effective, or that such a product is not therapeutically equivalent to other drug products. Rather, the exclusion may be based on the fact that FDA has not evaluated the safety, effectiveness, and quality of the drug product.

1.6 Practitioner/User Responsibilities

Professional care and judgment should be exercised in using the Orange Book. Evaluations of therapeutic equivalence for prescription drugs are based on scientific and medical evaluations by FDA. Products evaluated as therapeutically equivalent can be expected, in the judgment of FDA, to have the same clinical effect and safety profile when administered to patients under the conditions specified in the labeling. However, these products may differ in other characteristics that are not required by statute or regulation to be the same, such as shape, scoring configuration, release mechanisms, packaging, excipients (including colors, flavors, preservatives), expiration date/time, and, in some instances, labeling. If products with such differences are substituted for each other, there is a potential for patient confusion, e.g., due to differences in color or shape of tablets, inability to provide a given dose using a partial tablet if the proper scoring configuration is not available, or decreased patient acceptance of certain products because of flavor. There may also be patient-specific allergic reactions in rare cases due to a coloring or a preservative ingredient.

FDA evaluation of therapeutic equivalence in no way relieves practitioners of their professional responsibilities in prescribing and dispensing such products with due care and with appropriate information to individual patients. In those circumstances where the characteristics of a specific product, other than its active ingredient, are important in the therapy of a particular patient, the practitioner's prescribing of that product may be appropriate. Pharmacists must also be familiar with the different characteristics of therapeutically equivalent products, e.g., expiration dates/times and labeling directions for storage of the different products (particularly for reconstituted products), so they can properly advise patients when one product is substituted for another.

Multisource and single source drug products. In the Orange Book, FDA has evaluated for therapeutic equivalence only multisource prescription drug products approved under Section 505 of the FD&C Act, which in most instances means those pharmaceutical equivalents available (i.e., not on the Discontinued Drug Product list) from more than one manufacturer. For such products, a therapeutic equivalence code generally is included and product information is highlighted in bold face and underlined. Those products with approved applications that are single source (i.e., there is only one approved product available for that active ingredient, dosage form, route of administration, and strength) are also included in the Orange Book, but no therapeutic equivalence code is included with such products. Any drug product in the Orange Book repackaged and/or distributed by the applicant or some other person authorized by the applicant (e.g., an authorized generic) is considered to be therapeutically equivalent to the applicant's drug product, even if the applicant's drug product is single source or coded as non-equivalent (e.g., **BN**). Distributors or repackagers of an applicant's drug product are not identified in the Orange Book. The details of therapeutic equivalence codes and the policies underlying them are discussed in Section 1.7, *Therapeutic Equivalence Evaluations Codes*.

Products in the Orange Book are identified by the names of the holders of approved applications (applicants) who may not necessarily be the manufacturer of the product. There are numerous entities other than the applicant that may be involved in the development, manufacturing, and/or marketing of a product. Products listed in the Orange Book are identified by the applicant's name (firm name on the Form FDA 356h in the application). Where the applicant's name does not appear on the label, a person wishing to relate a specific product to the applicant name in the Orange Book may refer to FDA's NDC Directory¹⁶ and match its search terms to information on the label, such as the NDC Code if available.

Every product in the Orange Book is subject at all times to regulatory action. From time to time, approved products may be found in violation of one or more provisions of the FD&C Act. In such circumstances, the Agency may commence appropriate enforcement action to correct the violation, if necessary, by securing removal of the product from the market by voluntary recall, seizure, or other enforcement actions. Such regulatory actions are, however, independent of the inclusion of a product in the Orange Book. The main criterion for inclusion of a product is that it has an NDA or ANDA that has been approved and that has not been withdrawn for safety or effectiveness reasons. FDA believes that retention of a violative product in the Orange Book will not have any significant adverse health consequences, because other legal mechanisms are available to the Agency to prevent the product's actual marketing. FDA may, however, change a product's therapeutic equivalence rating if the circumstances giving rise to the violation change or otherwise call into question the Agency's assessment of whether a product meets the criteria for therapeutic equivalence.

1.7 Therapeutic Equivalence Evaluations Codes

Generally, prescription drug products that the Agency considers multisource have been assigned a therapeutic equivalence code. The coding system for therapeutic equivalence evaluations is designed to allow users to determine quickly whether the Agency has evaluated a particular approved prescription drug product (e.g., a particular strength of an approved drug that is not on the Discontinued Drug Product List) as therapeutically

¹⁶ <https://www.fda.gov/drugs/drug-approvals-and-databases/national-drug-code-directory>.

equivalent to other pharmaceutically equivalent prescription drug products (first letter) and to provide additional information on the basis of FDA's evaluations (second letter). With some exceptions (e.g., therapeutic equivalence evaluations for certain 505(b)(2) applications), the therapeutic equivalence evaluation date is the same as the approval date.

The two basic categories into which multisource drugs have been placed are indicated by the first letter of the relevant therapeutic equivalence code as follows:

A Drug products that FDA considers to be therapeutically equivalent to other pharmaceutically equivalent products, i.e., drug products for which:

- (1) there are no known or suspected bioequivalence problems. These are designated **AA, AN, AO, AP, or AT**, depending on the dosage form; or
- (2) actual or potential bioequivalence problems have been resolved with adequate *in vivo* and/or *in vitro* evidence supporting bioequivalence. These are designated **AB**.

B Drug products that FDA at this time, considers not to be therapeutically equivalent to other pharmaceutically equivalent products, i.e.,

drug products for which actual or potential bioequivalence problems have not been resolved by adequate evidence of bioequivalence. Often the problem is with specific dosage forms rather than with the active ingredients. These are designated **BC, BD, BE, BN, BP, BR, BS, BT, BX, or B***.

Individual drug products have been evaluated as therapeutically equivalent in accordance with the definitions and policies outlined below:

"A" CODES

Drug products that are considered to be therapeutically equivalent to other pharmaceutically equivalent products.

"A" products are those for which there are no known or suspected bioequivalence problems or for which actual or potential bioequivalence problems have been resolved with adequate *in vivo* and/or *in vitro* evidence supporting bioequivalence. Drug products designated with an "A" code fall under one of two main policies:

- (1) for those active ingredients or dosage forms for which no *in vivo* bioequivalence issue is known or suspected, the information necessary to show bioequivalence between pharmaceutically equivalent products is either presumed and considered self-evident (based on other information in the application for some dosage forms (e.g., solutions)), or satisfied by a showing that an acceptable *in vitro* approach is met. A therapeutically equivalent rating is assigned to such products so long as they are manufactured in accordance with Current Good Manufacturing Practice regulations and meet the other requirements of their approved applications (these are designated **AA, AN, AO, AP, or AT**, depending on the dosage form, as described below); or
- (2) for those Drug Efficacy Study Implementation (DESI) drug products containing active ingredients or dosage forms that have been identified by FDA as having actual or potential bioequivalence problems, and for post-1962 drug products presenting a potential bioequivalence problem,

an evaluation of therapeutic equivalence is assigned to pharmaceutical equivalents only if the approved application contains adequate scientific evidence establishing through *in vivo* and/or *in vitro* studies the bioequivalence of the product to a selected reference product (these products are designated as **AB**).

There are some general principles that may affect the substitution of pharmaceutically equivalent products in specific cases. Prescribers and dispensers of drugs should be alert to these principles so as to deal appropriately with situations that require professional judgment and discretion.

There may be labeling differences among pharmaceutically equivalent products that require attention on the part of the health professional (e.g., pharmaceutically equivalent powders to be reconstituted for administration as oral or injectable liquids may vary with respect to their expiration time or storage conditions after reconstitution). FDA's determination that such products are therapeutically equivalent is applicable only when each product is reconstituted, stored, and used under the conditions specified in its labeling.

The Agency may use notes in this publication to point out special situations, such as potential differences between two drug products that have been evaluated as bioequivalent and otherwise therapeutically equivalent, when they should be brought to the attention of health professionals. These notes are contained in Section 1.8, *Description of Certain Special Situations*. For example, in certain instances, there may be variations among therapeutically equivalent products in their use or in conditions of administration. When such variations may, in the Agency's opinion, affect prescribing or substitution decisions by health professionals, a note may be added to Section 1.8.

For example, occasionally a situation may arise in which changes in a listed drug after its approval (for example, a change in dosing interval) may have an impact on the substitutability of already approved generic versions of that product that were rated by the Agency as therapeutically equivalent to the listed drug. When such changes in the listed drug are considered by the Agency to have a significant impact on therapeutic equivalence, the Agency will change the therapeutic equivalence ratings for other versions of the drug product unless the manufacturers of those other versions of the product provide additional information to assure equivalence under the changed conditions. Pending receipt of the additional data, the Agency may add a note to Section 1.8, or, in rare cases, may even change the therapeutic equivalence rating.

In some cases (e.g., Isolyte® S w/ Dextrose 5% in Plastic Container and Plasma-Lyte® 148 and Dextrose 5% in Plastic Container), closely related products are listed as containing the same active ingredients, but in somewhat different amounts. In determining which of these products are pharmaceutically equivalent, generally the Agency has considered products to be pharmaceutically equivalent with labeled strengths of an ingredient that do not vary by more than 1%.

Different salts, esters or other noncovalent derivatives (such as a complex, chelate, or clathrate) of the same active moiety are regarded as different active ingredients. For the purpose of this publication, products containing such different active ingredients are considered pharmaceutical alternatives and, thus, are not therapeutically equivalent. Anhydrous and hydrated entities, as well as different polymorphs, are considered to be the

same active ingredient and are expected to meet the same standards for identity to be considered pharmaceutical equivalents and therapeutic equivalents.

The codes in this book are not intended to preclude healthcare professionals from converting pharmaceutically different concentrations into pharmaceutical equivalents using accepted professional practice.

Preservatives and other inactive ingredients may differ among some therapeutically equivalent drug products. These differences do not affect FDA's evaluation of therapeutic equivalence except in cases where these components may influence bioequivalence or routes of administration.

The specific sub-codes for those drugs evaluated as therapeutically equivalent and the policies underlying these sub-codes follow:

AA Products in conventional dosage forms not presenting bioequivalence problems

Multisource drug products coded as **AA** contain active ingredients and are in dosage forms that are not regarded as presenting either actual or potential bioequivalence problems or drug quality or standards issues. However, all oral dosage forms must, nonetheless, meet an appropriate *in vitro* bioequivalence standard that is acceptable to the Agency in order to be approved.

AB, AB1, AB2, AB3... Products meeting necessary bioequivalence requirements

Multisource drug products listed under the same heading (i.e., identical active ingredient(s), dosage form, and route(s) of administration) and having the same strength (see Section 1.2, *Therapeutic Equivalence-Related Terms, Strength*) generally will be coded **AB** if data and information are submitted demonstrating bioequivalence.

In certain instances, a number is added to the end of the **AB** code to make a three-character code (i.e., **AB1, AB2, AB3, etc.**). Three-character codes generally are assigned only in situations when more than one RLD of the same strength has been designated under the same heading. If a study is submitted that demonstrates bioequivalence to an RLD, the generic drug product will be given the same three-character code as the RLD it was compared against. For example, Adalat® CC and Procardia XL®, extended-release tablets, are listed under the active ingredient nifedipine. These drug products, listed under the same heading, are not bioequivalent to each other. Adalat® CC and Procardia XL® have been assigned ratings of **AB1** and **AB2**, respectively. Generic drug products deemed by FDA to be bioequivalent to Adalat® CC and Procardia XL® have been approved. As a result, the generic drug products bioequivalent to Adalat® CC have been assigned a rating of **AB1** and those bioequivalent to Procardia XL® have been assigned a rating of **AB2** (the assignment of an **AB1** or **AB2** rating to a specific product does not imply product preference). Even though drug products of distributors and/or repackagers are not included in the Orange Book, they are considered therapeutically equivalent to the applicant's drug product if the applicant's drug product is rated either with an **AB** or three-character code or is single source in the Orange Book. Drugs coded as **AB** under a heading are considered therapeutically equivalent only to other drugs coded as **AB** under that heading. Drugs coded with a three-character code under a heading are considered therapeutically equivalent only to other drugs coded with the same three-character code under that heading.

AN Solutions and powders for aerosolization

Uncertainty regarding the therapeutic equivalence of aerosolized products arises primarily because of differences in the drug delivery system. Solutions and powders intended for aerosolization that are marketed for use in general-use delivery systems are considered to be pharmaceutically and therapeutically equivalent and are coded **AN**. Those products that are compatible only with a specific delivery system or those products that are packaged in and with a specific delivery system are coded **BN**, unless they have met an appropriate bioequivalence standard and are otherwise determined to be therapeutically equivalent. Solutions or suspensions in a specific delivery system will be coded **AN** if the bioequivalence standard is based upon *in vitro* methodology. If bioequivalence needs to be demonstrated by *in vivo* methodology, then the drug products will be coded **AB**.

AO Injectable oil solutions

The absorption of drugs in injectable (parenteral) oil solutions may vary substantially with the type of oil employed as a vehicle and the concentration of the active ingredient. Injectable oil solutions are therefore considered to be pharmaceutically and therapeutically equivalent only when the active ingredient, its concentration, and the type of oil used as a vehicle are all identical.

AP Injectable aqueous solutions and, in certain instances, intravenous non-aqueous solutions

It should be noted that even though injectable (parenteral) products under a specific listing may be evaluated as therapeutically equivalent, there may be important differences among the products in the general category, Injectable; Injection. For example, historically some injectable products that are rated therapeutically equivalent are labeled for different routes of administration. In addition, some products evaluated as therapeutically equivalent may have different preservatives or no preservatives at all. Injectable products are available as solutions (e.g., concentrated sterile solutions for dilution or sterile solutions ready for injection) or powders for reconstitution (e.g., a lyophilized powder for injection). Solutions and powders for reconstitution are considered different dosage forms, and thus not pharmaceutical equivalent drug products, but may be considered pharmaceutical alternative drug products. Therefore, they are not rated as therapeutically equivalent (AP) to each other even if these pharmaceutical alternative drug products are designed to produce the same concentration prior to injection and are expected to have similar clinical effects and safety profile under the conditions of use described in the labeling. Consistent with accepted professional practice, it is the responsibility of the prescriber, dispenser, or individual administering the product to be familiar with a product's labeling to assure that it is given only by the route(s) of administration stated in the labeling. Historically, the dosage form of a parenteral product was listed as "injectable" while the product physically was generally either a solution or powder. Thus, different products listed with the historical term "injectable" do not necessarily have the same dosage form.

Certain commonly used large volume intravenous products in glass containers are not included in the Orange Book (e.g., dextrose injection 5%, dextrose injection 10%, sodium chloride injection 0.9%) since these

products are on the market without FDA approval and FDA has not published conditions for marketing such parenteral products under approved NDAs. When packaged in plastic containers, however, FDA regulations require approved applications prior to marketing. Approval then depends on, among other things, the extent of the available safety data involving the specific plastic component of the product. All large volume parenteral products are manufactured under similar standards, regardless of whether they are packaged in glass or plastic. Thus, FDA has no reason to believe that the packaging container of large volume parenteral drug products that are pharmaceutically equivalent would have any effect on their therapeutic equivalence.

Consistent with the definition of strength included in Section 1.2, *Therapeutic Equivalence-Related Terms*, the strength of a parenteral solution generally is identified by both the total drug content and the concentration of drug substance in a container approved by FDA. In the past, the strength of parenteral solutions in the Orange Book has not been fully displayed. Rather, the strength of parenteral solutions in the Orange Book has been displayed in terms of concentration, expressed as x mg/milliliter (mL).¹⁷ Generally, the amount of dry powder or lyophilized powder in a container is identified as the strength, expressed as x mg/vial.

However, FDA subsequently realized that the format of the Orange Book with respect to parenteral solutions should be changed to reflect that each strength of a drug is considered to be a separate listed drug. The Orange Book generally displays the strength of new approvals of parenteral solutions. Previously (i.e., prior to 2003), we would have displayed only the concentration of an approved parenteral solution, e.g., 1 mg/mL. For example, if this application had a 125 mL and 250 mL container approved, we would now display two product strengths, listing both total drug content and concentration of drug substance in the relevant approved container, e.g., 125 mg/125 mL (1 mg/mL) and 250 mg/250 mL (1 mg/mL).

AT Topical products

There are a variety of topical dosage forms available for dermatologic, ophthalmic, otic, rectal, and vaginal administration, including creams, gels, lotions, oils, ointments, pastes, solutions, sprays, suppositories, and inserts. Even though different topical dosage forms may contain the same active ingredient and potency, these dosage forms are not considered pharmaceutically equivalent. Therefore, they are not considered therapeutically equivalent. All solutions and DESI drug products containing the same active ingredient in the same topical dosage form for which a waiver of *in vivo* bioequivalence has been granted, or the application contains adequate scientific evidence establishing through an *in vitro* approach the bioequivalence of the product to a selected reference product, and for which chemistry and manufacturing processes are adequate to demonstrate bioequivalence, are considered therapeutically equivalent and coded **AT**. Pharmaceutically equivalent topical products that raise questions of bioequivalence and for which a waiver of *in vivo* bioequivalence has not been granted, including all post-1962 non-solution topical drug products, are coded **AB** when supported by adequate *in vivo* bioequivalence data, and **BT** in the absence of such data.

¹⁷ The concentrations of certain parenteral drug products, including contrast agents, may be expressed as a percentage rather than as x mg/mL.

"B" CODES

Drug products that FDA, at this time, considers not to be therapeutically equivalent to other pharmaceutically equivalent products.

"B" products, for which actual or potential bioequivalence problems have not been resolved by adequate evidence of bioequivalence, often have a problem with specific dosage forms rather than with the active ingredients. Drug products designated with a "B" code fall under one of three main policies:

- (1) the drug products contain active ingredients or are manufactured in dosage forms that have been identified by the Agency as having documented bioequivalence problems or a significant potential for such problems and for which no adequate studies demonstrating bioequivalence have been submitted to FDA; or
- (2) the quality standards are inadequate or FDA has an insufficient basis to determine therapeutic equivalence; or
- (3) the drug products are under regulatory review.

The specific coding definitions and policies for the "B" sub-codes are as follows:

B* Drug products requiring further FDA investigation and review to determine therapeutic equivalence

The code **B*** is assigned to products previously assigned an **A** or **B** code when FDA receives new information that raises a significant question regarding therapeutic equivalence that can be resolved only through further Agency investigation and/or review of data and information submitted by the applicant. The **B*** code signifies that the Agency will take no position regarding the therapeutic equivalence of the product until the Agency completes its investigation and review.

BC Extended-release dosage forms (capsules, injectables and tablets)

Extended-release tablets are formulated in such a manner as to make the contained drug substance available over an extended period of time following ingestion.

Although bioavailability studies have been conducted on these dosage forms, they may be subject to bioavailability differences, primarily because applicants developing extended-release products for the same active ingredient rarely employ the same formulation approach. FDA, therefore, does not consider different extended-release dosage forms containing the same active ingredient in equal strength to be therapeutically equivalent unless equivalence between individual products in both rate and extent has been specifically demonstrated through appropriate bioequivalence studies. Extended-release products for which such bioequivalence data have not been submitted are coded **BC**, while those for which such data are available have been coded **AB**.

BD Active ingredients and dosage forms with documented bioequivalence problems

The **BD** code denotes products containing active ingredients with known bioequivalence problems and for which adequate studies have not been

submitted to FDA demonstrating bioequivalence. Where studies showing bioequivalence have been submitted, the product has been coded **AB**.

BE Delayed-release oral dosage forms

Where the drug may be destroyed or inactivated by the gastric juice or where it may irritate the gastric mucosa, the use of "enteric" coatings is indicated. Such coatings are intended to delay the release of the medication until the tablet has passed through the stomach. Drug products in delayed-release dosage forms containing the same active ingredients may be subject to significant differences in absorption. Unless otherwise specifically noted, the Agency considers different delayed-release products containing the same active ingredients as presenting a potential bioequivalence problem and codes these products **BE** in the absence of *in vivo* studies showing bioequivalence. If adequate *in vivo* studies have demonstrated the bioequivalence of specific delayed-release products, such products are coded **AB**.

BN Products in aerosol-nebulizer drug delivery systems

This code applies to drug solutions or powders that are marketed only as a component of, or as compatible with, a specific drug delivery system. There may, for example, be significant differences in the dose of drug and particle size delivered by different products of this type. Therefore, the Agency does not consider different metered aerosol dosage forms containing the same active ingredient(s) in equal strengths to be therapeutically equivalent unless the drug products meet an appropriate bioequivalence standard; such products are coded **AB**.

BP Active ingredients and dosage forms with potential bioequivalence problems

FDA's bioequivalence regulations (21 CFR 320.33) contain criteria and procedures for determining whether a specific active ingredient in a specific dosage form has a potential for causing a bioequivalence problem. It is FDA's policy to consider an active ingredient meeting these criteria as having a potential bioequivalence problem even in the absence of positive data demonstrating inequivalence. Pharmaceutically equivalent products containing these active ingredients in oral dosage forms are coded **BP** until adequate bioequivalence data are submitted, after which such products are coded **AB**. Injectable suspensions containing an active ingredient suspended in an aqueous or oleaginous vehicle have also been coded **BP**. Injectable suspensions are subject to bioequivalence problems because differences in particle size, polymorphic structure of the suspended active ingredient, or the suspension formulation can significantly affect the rate of release and absorption. FDA does not consider pharmaceutical equivalents of these products bioequivalent without adequate evidence of bioequivalence; such products would be coded **AB**.

BR Suppositories or enemas that deliver drugs for systemic absorption

The absorption of active ingredients from suppositories or enemas that are intended to have a systemic effect (as distinct from suppositories administered for local effect) can vary significantly from product to product. Therefore, FDA considers pharmaceutically equivalent systemic suppositories or enemas bioequivalent only if *in vivo* evidence of bioequivalence is available. In those cases where *in vivo* evidence is

available, the products are coded **AB**. If such evidence is not available, the products are coded **BR**.

BS Products having drug standard deficiencies

If the drug standards for an active ingredient in a particular dosage form are found by FDA to be deficient so as to prevent an FDA evaluation of either pharmaceutical or therapeutic equivalence, all drug products containing that active ingredient in that dosage form are coded **BS**. For example, if the standards permit a wide variation in pharmacologically active components of the active ingredient such that pharmaceutical equivalence is in question, all products containing that active ingredient in that dosage form are coded **BS**.

BT Topical products with bioequivalence issues

This code applies mainly to post-1962 dermatologic, ophthalmic, otic, rectal, and vaginal products for topical administration, including creams, gels, lotions, oils, ointments, pastes, solutions, sprays, suppositories, and inserts not intended for systemic drug absorption. Topical products evaluated as having acceptable clinical performance, but that are not bioequivalent to other pharmaceutically equivalent products or that lack sufficient evidence of bioequivalence, will be coded **BT**.

BX Drug products for which the data are insufficient to determine therapeutic equivalence

The code **BX** is assigned to specific drug products for which the data that have been reviewed by the Agency are insufficient to determine therapeutic equivalence under the policies stated in this document. In these situations, the drug products are presumed to be therapeutically inequivalent until the Agency has determined that there is adequate information to make a full evaluation of therapeutic equivalence.

1.8 Description of Certain Special Situations

Certain drugs listed in the Orange Book present special situations that merit further discussion. The following are descriptions of certain examples of those special situations:

Amino Acid and Protein Hydrolysate Injections. These products differ in the amount and kinds of amino acids they contain and, therefore, are not considered pharmaceutical equivalents. For this reason, these products are not considered therapeutically equivalent. At the same time, the Agency believes that it is appropriate to point out that where nitrogen balance is the sole therapeutic objective and individual amino acid content is not a consideration, pharmaceutical alternatives with the same total amount of nitrogen content may be expected to have the same clinical effect and safety profile when administered to patients under the conditions specified in the labeling.

Gaviscon®. Gaviscon® is an OTC product that has been marketed since September 1970. The active ingredients in this product, aluminum hydroxide and magnesium trisilicate, were reviewed by the Agency's OTC Antacid Panel and were considered to be safe and effective ingredients (Category I) by that Panel. However, the tablet failed to pass the antacid test that is required of all antacid products. The Agency, therefore, placed the tablet in Category III for lack of effectiveness. A full NDA with clinical studies was

submitted by Marion Laboratories, Inc., and approved by FDA on December 9, 1983. Gaviscon®'s activity in treating reflux acidity is made possible by the physical-chemical properties of the inactive ingredients, sodium bicarbonate and alginic acid. Therefore, *all ANDAs that cite Gaviscon® tablets as the RLD must contain the inactive ingredients sodium bicarbonate and alginic acid.* A full NDA will be required to support the effectiveness of the drug product if different inactive ingredients are to be substituted for sodium bicarbonate or alginic acid or if different proportions of these ingredients are to be used.

Levothyroxine Sodium.¹⁸ Because there are multiple RLDs for levothyroxine sodium tablets and some RLD applicants have conducted studies to establish their drugs' therapeutic equivalence to other RLDs, FDA has determined that its usual practice of assigning two- or three-character therapeutic equivalence codes may be potentially confusing and inadequate for these drug products. Looking at the Orange Book listing alone for a product identified as an RLD or reference standard, it may be difficult to determine to which therapeutic equivalence code the RLDs and/or reference standard designation corresponds. For example, Unithroid 0.3 mg strength has been assigned the therapeutic equivalence codes AB1, AB2, and AB3 and it is identified as the RLD and reference standard, but it is unclear that the RLD and reference standard designations are associated with the AB1 therapeutic equivalence code.

Accordingly, FDA provides the following chart, which identifies (1) an RLD for each therapeutic equivalence code in the Orange Book and (2) and the reference standard products in the Active Section of the Orange Book.¹⁹

- Therapeutic equivalence has been established between products that have the same AB+number therapeutic equivalence code (i.e., AB1, AB2, AB3 or AB4).
- More than one therapeutic equivalence code may apply to some products. One common therapeutic equivalence code indicates therapeutic equivalence between products. For example, Unithroid has been assigned therapeutic equivalence codes AB1, AB2, and AB3, and therefore, Unithroid tablets are considered therapeutically equivalent to other levothyroxine sodium products of the same strength with these therapeutic equivalence codes.

TE Code	Proprietary Name	Applicant	Strength	Appl No	RLD	RS
AB1	UNITHROID	STEVENS J	0.3 MG	N021210	RLD	RS
AB2	SYNTHROID	ABBVIE	0.3 MG	N021402	RLD	RS
AB3	LEVOXYL	KING PHARMS	0.2 MG	N021301	RLD	RS

¹⁸ In previous editions of the Orange Book, FDA provided a chart outlining therapeutic equivalence codes for all 0.025 mg levothyroxine sodium drug products in the Active Section of the Orange Book. FDA has decided, for ease of review, to revise the chart to identify the NDAs for the reference listed drugs for each therapeutic equivalence code (i.e., AB1, AB2, AB3, and AB4), and their corresponding reference standards, which are identified in 0.2 and 0.3 mg strengths.

¹⁹ The chart is current as of the date of publication of the annual edition. See the most current monthly cumulative supplement for updates to this information available at <https://www.fda.gov/media/72973/download>. Please consult the Active Section for information on other strengths.

AB4	THYRO-TABS	ALVOGEN INC	0.3 MG	N021116	RLD	-
AB4	LEVOTHYROXINE SODIUM ²⁰	MYLAN	0.3 MG	A076187	-	RS

Patent Certification(s) and Reference Standard for ANDAs Duplicating a Drug Product Approved in a Petitioned ANDA. To submit an ANDA for a generic drug that is not the same as its RLD because it has one different active ingredient in a fixed-combination drug product, or has a different route of administration, dosage form, or strength than that of the RLD, an applicant first must obtain permission from FDA through what is known as a suitability petition pursuant to Section 505(j)(2)(C) of the FD&C Act. A petitioned ANDA relies on the RLD described in the suitability petition. An ANDA seeking approval of a drug that is the same as a drug product approved in a petitioned ANDA should use as its RLD, the RLD that served as the basis for the approved suitability petition, and use the drug product approved in the petitioned ANDA as its reference standard for conducting an *in vivo* bioequivalence study required for approval. However, the RLD for any such ANDA is generally the listed drug referenced in the approved suitability petition. The ANDA must include appropriate patent certification(s) and an exclusivity statement with respect to the RLD that served as the basis for the approved suitability petition²¹ (This concept also generally applies to an ANDA applicant that utilizes a reference standard that is not an RLD, as such an application must include appropriate patent certification(s) and an exclusivity statement with respect to the RLD).

Waived exclusivity. If an NDA submitted under Section 505(b) of the FD&C Act qualifies for exclusivity under the FD&C Act, the exclusivity is generally listed in the Patent and Exclusivity Information Addendum of the Orange Book. If a drug product has qualified for this exclusivity, FDA will not accept for review and/or will not approve, as applicable, other applications blocked by the relevant exclusivity. If the listed drug is also protected by one or more patents, the approval date for an ANDA or 505(b)(2) application that relies on the listed drug will be determined based on an analysis of the applicant's patent certification(s) or statement(s) for each relevant patent and the effect of relevant exclusivity listed in the Orange Book. However, the holder of the NDA may waive its exclusivity as to any or all applications that might otherwise be blocked by such exclusivity. If an NDA holder waives its exclusivity, qualified applications may be accepted for review and/or approved, as applicable. An NDA for which the holder has waived its exclusivity as to all applications will be coded with a "W" in the Patent and Exclusivity Section of the Orange Book. The applicant whose

²⁰ Alvogen, Inc.'s tablets (NDA 021116) (previously known as Levothroid) previously was listed in the Discontinued Drug Product List section of the Orange Book. It is the RLD for therapeutic equivalents identified with the AB4 code. During this time, Mylan's levothyroxine product (ANDA 076187) was selected as the reference standard for ANDA applicants to use to establish bioequivalence to Thyro-Tabs. It remains the reference standard for ANDA applicants to use to establish bioequivalence to Thyro-Tabs. If an ANDA that uses Mylan's levothyroxine product as its reference standard is approved, the ANDA will receive an AB4 rating. The ANDA applicant also may obtain an AB rating for its product to the other RLDs (i.e., Unithroid, Synthroid, and Levoxyl) by submitting supplements that demonstrate that the generic product is bioequivalent to these other RLDs and satisfies all other therapeutic equivalence criteria with respect to these reference listed drugs. See Letter from Janet Woodcock, M.D., Director, Center for Drug Evaluation and Research, FDA to Teri Nataline, Principal Consultant, Lachman Consultant Services, Inc., Docket No. FDA-2015-P-0403 (May 27, 2016).

²¹ If after approval of a suitability petition and before approval of an ANDA submitted pursuant to the approved petition, a drug product is approved in an NDA for the change described in the petition, the suitability petition and the RLD identified in the petition can no longer be the basis of submission for such ANDA. Under these circumstances, an applicant seeking approval for a drug product with the change approved in the suitability petition must submit a new ANDA that identifies the drug product approved under such NDA as the RLD and comply with applicable regulatory requirements. See 21 CFR 314.93(f)(2).

product might otherwise be blocked by this exclusivity should indicate in the exclusivity statement in its application that the holder of the listed drug has waived its exclusivity.

1.9 Therapeutic Equivalence Code Change for a Category of Multisource Drug Products

The Agency will use the following procedures when, in response to a petition or on its own initiative, it is considering a change in the therapeutic equivalence code for approved multisource drug products. Such changes will generally occur when the Agency becomes aware of new scientific information affecting the therapeutic equivalence of an entire category of multisource drug products in the Orange Book (e.g., information concerning the active ingredient or the dosage form), rather than information concerning a single drug product within the category. These procedures will be used when a change in therapeutic equivalence code is under consideration for all drug products found in the Prescription Drug Product List under a specific active ingredient and dosage form. The change may be from the code signifying that the drug does not present a bioequivalence problem (e.g., **AA**) to a code signifying an actual or potential bioequivalence problem (e.g., **BP**), or vice versa. This procedure does not apply to a change of a particular product code (e.g., a change from **BP** to **AB** or from **AB** to **BX**).

Before making a change in a therapeutic equivalence code for an entire category of multisource drug products as described above, the Agency will announce in the *Introduction* to the Cumulative Supplement that it is considering the change and will invite comments. Comments, along with scientific data, may be sent to the Director, Office of Bioequivalence, Office of Generic Drugs, Center for Drug Evaluation and Research, Food and Drug Administration, Central Document Room, 5901-B Ammendale Rd., Beltsville, MD 20705-1266.

The comment period will generally be 60 days in length, and the closing date for comments will be listed in the description of the proposed change for each drug entity.

The most useful type of scientific data submitted to support comments is generally an *in vivo* bioavailability/bioequivalence study conducted on batches of the subject drug products. Comments including scientific data from an *in vivo* bioavailability/bioequivalence study should present a full description of the analytical procedures and equipment used, a validation of the analytical methodology, including the standard curve, a description of the method of calculating results, and a description of the pharmacokinetic and statistical models used in analyzing the data. Anecdotal or testimonial information is the least useful to the Agency, and submission of comments based on such information is discouraged. However, when there is supporting published or unpublished scientific literature, copies should be submitted with comments.

1.10 Change of the Therapeutic Equivalence Evaluation for a Single Product

The procedure described in Section 1.9 does not apply to a change in a single drug product code. For example, a change in a single drug product's code from **BP** to **AB** as a result of the submission of an acceptable bioequivalence study ordinarily will not be the subject of notice and comment in the Cumulative Supplement. Likewise, a change in a single drug product's code from **AB** to **BX** (e.g., as a result of new information raising a significant question as to bioequivalence) does not require notice and comment. The Agency's responsibility to provide the public with the Agency's

most current information related to therapeutic equivalence may require a change in a drug product's code prior to any formal notice and opportunity for the applicant to be heard. The publication in the *Federal Register* of a proposal to withdraw approval of a drug product will ordinarily result in a change in a product's code from **AB** to **BX** if this action has not already been taken.

We recognize that certain drug products approved in 505(b)(2) applications might not have therapeutic equivalence codes, and that FDA may undertake therapeutic equivalence evaluations with respect to such drug products. A person seeking to have a therapeutic equivalence rating for a drug product approved in a 505(b)(2) application may petition the Agency through the citizen petition procedure (see 21 CFR 10.25(a) and 21 CFR 10.30).²²

1.11 Discontinued Drug Product List

The drug products in the Discontinued Drug Product List of the Orange Book (Discontinued Drug Product List) for which a determination has been made that the products were not withdrawn for safety or effectiveness reasons have been annotated with a footnote following the product strength: "***Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons***". The determinations listed in the Orange Book are only reflective of determinations made since 1995 and published in the *Federal Register*. The identification of these drug products in the Discontinued Drug Product List should avoid the submission of multiple citizen petitions requesting a determination for the same drug product.

Generally, approved products are added to the Discontinued Drug Product List when the applicant notifies the Division of Orange Book Publication and Regulatory Assessment (DOBPR) of the products' not-marketed status. Products may also be added to the Discontinued Drug Product List if annual reports or other submissions to the Agency indicate the product is not being marketed or as a result of other Agency administrative actions.²³ Changes to the Orange Book are not affected by the drug registration and listing requirements of Section 510 of the FD&C Act.

1.12 Changes to the Orange Book

Every effort is made to ensure the Annual Edition is current and accurate. Applicants are requested to inform DOBPR of any changes or corrections, including any change in ownership or a product's marketing status that would result in the product being moved to the Discontinued Drug Product List. FDA notes that under Section 506I(a) of the FD&C Act, application holders must notify the Agency in writing 180 days prior to withdrawing a drug product from sale, or if 180 days is not practicable, not later than the date of withdrawal from sale. Furthermore, Section 506I(b) of the FD&C Act requires that application holders notify the Agency in writing within 180 days of approval of a drug product if such drug product will not be available for sale within 180 days of approval. A request to include a newly approved product in the Discontinued Drug Product List, rather than parts 1 or 2 of the Orange Book (as discussed in Section 1.1), must be submitted to DOBPR by the end of the month in which the product is approved

²² Section 3222 of the Food and Drug Omnibus Reform Act of 2022 (enacted December 29, 2022) amended the FD&C Act by adding a new provision to Section 505(j)(7)(A). Section 505(j)(7)(A)(v)(I) sets forth certain conditions under which FDA considers therapeutic equivalence evaluation requests in an application for an eligible drug submitted or approved pursuant to Section 505(b)(2) of the FD&C Act.

²³ See, e.g., Section 506I(d) of the FD&C Act.

to ensure that the product is not included in the Active Section of the next published Orange Book update. The Prescription Drug Product List and the OTC Drug Product List are collectively referred to as the "Active Section."

In addition, DOBPRA generally will act on requests to change a proprietary name for a listed drug only after approval of a supplement for the relevant change in proprietary name. To the extent that conventions for describing product identification information (i.e., active ingredients, dosage forms, routes of administration, product names, applicants, strengths) evolve over time, the Agency generally does not intend to revise such information for drug products already listed in the Orange Book, but rather intends to apply the change prospectively to drug products as they are added to the Orange Book.

You can contact DOBPRA by email at orangebook@fda.hhs.gov.

1.13 Availability of the Edition

The Annual Edition and current monthly Cumulative Supplement are available in a Portable Document Format (PDF) at the [Orange Book](#) home page by clicking on Publications.

2.0 HOW TO USE THE DRUG PRODUCT LISTS

2.1 Key Sections for Using the Drug Product Lists

This publication contains illustrations, along with Drug Product Lists, indices, and lists of abbreviations and terms which facilitate their use.

Illustrations. The annotated *Drug Product Illustration* (see Section 2.2) and the *Therapeutic Equivalence Evaluations Illustration* (see Section 2.3) are offered to provide further clarification. These depict the format found in the Prescription Drug Product List (the only list in which therapeutic equivalence evaluation codes are displayed).

Drug Product Lists. The Prescription and OTC Drug Product Lists, arranged alphabetically by active ingredient(s), contain product identification information (active ingredients, dosage forms, routes of administration, product names, applicants, strengths) for single and multiple ingredient drug products. Also shown are the application number and drug product number (FDA internal computer data use only) and approval dates for those drug products approved on or after January 1, 1982. The application number preceded by "N" is a New Drug Application (NDA or commonly the innovator). The application number preceded by an "A" is an Abbreviated New Drug Application (ANDA or commonly the generic).

The Discontinued Drug Product List, arranged alphabetically by active ingredient(s), contains product identification information (active ingredients, dosage forms, routes of administration, product names, applicants, strengths).

If a prescription drug product is available from more than one source (multisource), a therapeutic equivalence code will appear in front of the applicant's name. If a product is therapeutically equivalent to one or more products or to an appropriate reference, it will be designated with a code beginning with "A" and the entry will be underlined and printed in bold font for emphasis.

Active ingredient headings for multiple ingredient (combination) drug products are arranged alphabetically. For purposes of this publication, this alphabetical sort takes precedence over United States Pharmacopeia official monograph order (i.e., Reserpine, Hydralazine Hydrochloride, Hydrochlorothiazide). For example, product information labeled as Reserpine, Hydrochlorothiazide and Hydralazine Hydrochloride appears under the active ingredient heading *Hydralazine Hydrochloride; Hydrochlorothiazide; Reserpine*. A cross-reference to the product information (for prescription and OTC products) appears for each additional active ingredient in the product. For combination drug products, the ingredient strengths are separated by semicolons and appear in the same relative sequence as the ingredients in the heading. Available strengths of the dosage form from an applicant appear on separate lines.

To use the Drug Product Lists, determine by alphabetical order the ingredient under which the product information is listed, using the Product Name Index, if necessary. Then, find the ingredient in the applicable Drug Product List. Proceed to the dosage form and route of administration and

compare products within that ingredient heading only. Therapeutic equivalence or inequivalence for prescription products is determined on the basis of the therapeutic equivalence codes provided within that specific dosage form and route heading. The OTC Drug Product List, Discontinued Drug Product List, and Drug Products with Approval under Section 505 of the Act Administered by the Center for Biologics Evaluation and Research List have their data arranged similarly.

The Discontinued Drug Product List contains approved products that have never been marketed, have been discontinued from marketing and we have not determined that they were withdrawn for safety or effectiveness reasons, are for military use, or have had their approvals withdrawn for other than safety or efficacy reasons subsequent to being discontinued from marketing. All products having a "@" in the December Cumulative Supplement of the previous Edition List have been added to the Discontinued Drug Product List appearing in this Edition. In addition, approved drug products that are not in the commercial distribution channel e.g., approved drug products in applications for export only are also listed in the Discontinued Drug Product List.

Product Name Index (*Prescription and OTC Drug Product Lists*). This is an index of drug products by trade name or established name of the active ingredient, if no trade name exists. The second term of each entry indicates the active ingredient name under which product information can be found in the appropriate Drug Product List. For those drug products with multiple active ingredients, only the first active ingredient (in alphabetical order) will appear. OTC products are so designated.

Product Name Index Listed by Applicant (*Prescription and OTC Drug Product Lists*). This is an index that cross-references applicants to drug products. The bolded and underlined entry represents the applicant name abbreviation used in this publication. Each complete applicant name that is represented by the abbreviated name is marked with an asterisk (*). Listed under each complete applicant name is the first alphabetically arranged ingredient under which product information can be found in the appropriate Drug Product List. OTC products are so designated. To use the Drug Product Lists, determine by alphabetical order the ingredient under which the product information is listed, using the Product Name Index, if appropriate.

Uniform Terms. To improve readability, uniform terms are used to designate dosage forms, routes of administration, and abbreviations used to express strengths. These terms are listed in Appendix C. In some cases, the terms used may differ from those used in product labels and other labeling.

2.2 DRUG PRODUCT ILLUSTRATION

SINGLE INGREDIENT

ACTIVE INGREDIENT	→	<u>MEPERIDINE HYDROCHLORIDE</u>						
DOSAGE FORM; ROUTE OF ADMINISTRATION	→	INJECTABLE; INJECTION						
TRADE OR GENERIC NAMES	→	<u>HEXANON</u>						
REFERENCE LISTED DRUG* (+)	→	<u>AP</u> +!	PAGE PHARMA	<u>25MG/ML</u>	<u>N013111</u>	<u>001</u>	AUG 22,	1983
REFERENCE STANDARD * (!)	→	<u>AP</u> +!		<u>50MG/ML</u>	<u>N013111</u>	<u>002</u>	AUG 22,	1983
	→	<u>AP</u> +!		<u>75MG/ML</u>	<u>N013111</u>	<u>003</u>	AUG 22,	1983
	→	<u>AP</u> +!		<u>100MG/ML</u>	<u>N013111</u>	<u>004</u>	JAN 04,	1989
	→	<u>MEPERIDINE HCL</u>						
	→	<u>AP</u>	DAVIS PHARM	<u>25MG/ML</u>	<u>A064890</u>	<u>001</u>	FEB 29,	1987
	→	<u>AP</u>		<u>50MG/ML</u>	<u>A064890</u>	<u>002</u>	FEB 29,	1987
	→	<u>AP</u>		<u>75MG/ML</u>	<u>A064890</u>	<u>003</u>	FEB 29,	1987
	→	<u>AP</u>		<u>100MG/ML</u>	<u>A064890</u>	<u>004</u>	MAR 08,	1992
THERAPEUTIC EQUIVALENCE (TE) CODE FOR MULTISOURCE PRODUCT	→							
SINGLE SOURCE PRODUCT (NO TE CODE)	→	<u>AP</u>	! TIMOKIM LLC	10MG/ML	A099225	001	DEC 12,	1995
	→	<u>AP</u>	JOHNSON MED	<u>25MG/ML</u>	<u>A099226</u>	<u>001</u>	NOV 27,	1993
	→	<u>AP</u>	! KENDRA PHARM	150MG/ML	A079444	001	OCT 31,	1999
APPLICANT	→							
AVAILABLE STRENGTH(S) OF A PRODUCT	→							
APPLICATION NUMBER	→							
PRODUCT NUMBER IS FOR FDA INTERNAL COMPUTER DATA USE ONLY	→							
APPROVAL DATE	→							

*NOTE: REFERENCE LISTED DRUG AND REFERENCE STANDARD ARE DISCUSSED IN THE PREFACE SECTION 1.4

MULTIPLE INGREDIENTS WITH PRODUCT INFORMATION

ALPHABETICALLY SORTED BY ACTIVE INGREDIENT	→	<u>HYDRALAZINE HYDROCHLORIDE; HYDROCHLOROTHIAZIDE; RESERPINE</u>								
PRODUCT INFORMATION	→	<table border="0"> <tr> <td>TABLET; ORAL</td> <td></td> </tr> <tr> <td>HYDROCHLOROTHIAZIDE, RESERPINE AND HYDRALAZINE HCL</td> <td></td> </tr> <tr> <td>REINWALD LABS</td> <td>25MG; 15MG; 0.1MG</td> </tr> <tr> <td>A069808</td> <td>001 JAN 18, 1982</td> </tr> </table>	TABLET; ORAL		HYDROCHLOROTHIAZIDE, RESERPINE AND HYDRALAZINE HCL		REINWALD LABS	25MG; 15MG; 0.1MG	A069808	001 JAN 18, 1982
TABLET; ORAL										
HYDROCHLOROTHIAZIDE, RESERPINE AND HYDRALAZINE HCL										
REINWALD LABS	25MG; 15MG; 0.1MG									
A069808	001 JAN 18, 1982									

THIS EXAMPLE IS FOR PURPOSE OF ILLUSTRATION ONLY. IT DOES NOT REPRESENT ACTUAL PRODUCTS FROM THE PRESCRIPTION DRUG PRODUCT LIST.

2.3 THERAPEUTIC EQUIVALENCE EVALUATIONS ILLUSTRATION

DRUG PRODUCTS CODED **AB** (OR ANY CODE BEGINNING WITH AN "A") UNDER AN INGREDIENT AND DOSAGE FORM HEADING ARE CONSIDERED THERAPEUTICALLY EQUIVALENT ONLY TO OTHER PRODUCTS CODED **AB** (OR ANY CODE BEGINNING WITH AN "A") AND **NOT** TO THOSE CODED **BP** (OR ANY CODE BEGINNING WITH "B") AND ANY PRODUCTS NOT LISTED. DRUG PRODUCTS CODED **BP** (OR ANY CODE BEGINNING WITH A "B") ARE **NOT** CONSIDERED THERAPEUTICALLY EQUIVALENT TO ANY OTHER PRODUCT. FOR A COMPLETE EXPLANATION OF THE **TE** CODES REFER TO SECTION 1.7 OF THE *INTRODUCTION*.

SULFASALAZINE

TABLET; ORAL

FAZINE

AB PARKLAND **500MG** **A042999** **001**

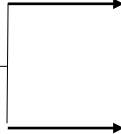
SULAZINE

AB URSA **500MG** **A042222** **001**

SULFASALAZINE

BP BROWN 500MG A041297 001

PRODUCTS CONSIDERED THERAPEUTICALLY EQUIVALENT TO EACH OTHER



PRODUCTS CONSIDERED **NOT** THERAPEUTICALLY EQUIVALENT TO ANY OTHER PRODUCTS LISTED

SULFASALAZINE

TABLET; ORAL

FAZINE

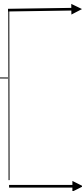
AB PARKLAND **500MG** **A042999** **001**

SULFASALAZINE

BP BROWN 500MG A041297 001

SOUTH 500MG A067627 001

PRODUCTS CONSIDERED **NOT** THERAPEUTICALLY EQUIVALENT TO EACH OTHER



NOTE: BOLD FONT AND UNDERLINING DENOTES MULTISOURCE PRODUCTS WHICH ARE CONSIDERED THERAPEUTICALLY EQUIVALENT.

THIS EXAMPLE IS FOR PURPOSES OF ILLUSTRATION ONLY. IT DOES NOT REPRESENT ACTUAL PRODUCTS FROM THE PRESCRIPTION DRUG PRODUCT LIST.

PRESCRIPTION DRUG PRODUCT LIST

ABACAVIR SULFATE

SOLUTION;ORAL

ABACAVIR SULFATE

AA	AUROBINDO PHARMA LTD	EQ 20MG BASE/ML	A077950 001	Mar 14, 2018
AA	HETERO LABS LTD III	EQ 20MG BASE/ML	A201107 001	Sep 26, 2016

ZIAGEN

AA	+! VIIV HLHCARE	EQ 20MG BASE/ML	N020978 001	Dec 17, 1998
-----------	-----------------	------------------------	--------------------	--------------

TABLET;ORAL

ABACAVIR SULFATE

AB	AUROBINDO PHARMA LTD	EQ 300MG BASE	A077844 001	Dec 17, 2012
AB	CHARTWELL RX	EQ 300MG BASE	A091050 001	Oct 28, 2016
AB	CIPLA	EQ 300MG BASE	A078119 001	Nov 21, 2017
AB	! HETERO LABS LTD III	EQ 300MG BASE	A091560 001	Sep 13, 2013
AB	MYLAN PHARMS INC	EQ 300MG BASE	A091294 001	Jun 18, 2012

ABACAVIR SULFATE; DOLUTEGRAVIR SODIUM; LAMIVUDINE

TABLET;ORAL

TRIUMEQ

+!	VIIV HLHCARE	EQ 600MG BASE;EQ 50MG BASE;300MG	N205551 001	Aug 22, 2014
----	--------------	----------------------------------	-------------	--------------

TABLET, FOR SUSPENSION;ORAL

TRIUMEQ PD

+!	VIIV HLHCARE	EQ 60MG BASE;EQ 5MG BASE;30MG	N215413 001	Mar 30, 2022
----	--------------	-------------------------------	-------------	--------------

ABACAVIR SULFATE; LAMIVUDINE

TABLET;ORAL

ABACAVIR SULFATE AND LAMIVUDINE

AB	AUROBINDO PHARMA LTD	EQ 600MG BASE;300MG	A090159 001	Nov 15, 2018
AB	CHARTWELL RX	EQ 600MG BASE;300MG	A204990 001	Mar 28, 2017
AB	! CIPLA	EQ 600MG BASE;300MG	A091144 001	Mar 28, 2017
AB	LAURUS	EQ 600MG BASE;300MG	A216332 001	Jul 25, 2022
AB	MACLEODS PHARMS LTD	EQ 600MG BASE;300MG	A212663 001	Dec 19, 2024

ABACAVIR SULFATE; LAMIVUDINE; ZIDOVUDINE

TABLET;ORAL

ABACAVIR SULFATE, LAMIVUDINE AND ZIDOVUDINE

!	LUPIN LTD	EQ 300MG BASE;150MG;300MG	A202912 001	Dec 05, 2013
---	-----------	---------------------------	-------------	--------------

ABALOPARATIDE

SOLUTION;SUBCUTANEOUS

TYMLOS

+!	RADIUS	3.12MG/1.56ML (2MG/ML)	N208743 001	Apr 28, 2017
----	--------	------------------------	-------------	--------------

ABEMACICLIB

TABLET;ORAL

VERZENIO

+	ELI LILLY AND CO	50MG	N208716 001	Sep 28, 2017
+		100MG	N208716 002	Sep 28, 2017
+		150MG	N208716 003	Sep 28, 2017
+!		200MG	N208716 004	Sep 28, 2017

ABIRATERONE ACETATE

TABLET;ORAL

ABIRATERONE ACETATE

AB	AMNEAL PHARMS	250MG	A208327 001	Jan 07, 2019
AB		500MG	A208327 002	Dec 23, 2020
AB	APOTEX	250MG	A208453 001	Oct 31, 2018
AB	DR REDDYS	250MG	A208416 001	May 18, 2020
AB		500MG	A208416 002	Sep 01, 2023
AB	FLORIDA	500MG	A215086 001	Mar 23, 2023
AB	GLENMARK SPECLT	250MG	A209227 001	Oct 16, 2019
AB		500MG	A209227 002	May 19, 2022
AB	HIKMA	250MG	A208339 001	Oct 31, 2018
AB	MSN	250MG	A210686 001	Jul 10, 2019
AB		500MG	A210686 002	Apr 24, 2024
AB	MYLAN	250MG	A208446 001	Oct 31, 2018
AB		500MG	A208446 002	Dec 14, 2020
AB	NOVUGEN	250MG	A215947 001	Jan 05, 2022
AB		500MG	A215947 002	Jan 05, 2022
AB	QILU	250MG	A212462 001	Sep 27, 2019
AB		500MG	A212462 002	Jun 25, 2021
AB	RISING	250MG	A208371 001	Feb 25, 2019
AB	TEVA PHARMS USA	250MG	A208432 001	Oct 31, 2018
AB		500MG	A210726 001	Jan 26, 2023

PRESCRIPTION DRUG PRODUCT LIST

ABIRATERONE ACETATE

TABLET; ORAL

ABIRATERONE ACETATE

<u>AB</u>	WOCKHARDT BIO AG	<u>250MG</u>	<u>A208380</u>	<u>001</u>	Feb 27, 2019
<u>ZYTIGA</u>					
<u>AB</u>	+	JANSSEN BIOTECH	<u>250MG</u>	<u>N202379</u>	<u>001</u> Apr 28, 2011
<u>AB</u>	+	!	<u>500MG</u>	<u>N202379</u>	<u>002</u> Apr 14, 2017
YONSA					
	+	SUN PHARM	125MG	N210308	001 May 22, 2018

ABIRATERONE ACETATE; NIRAPARIB TOSYLATE

TABLET; ORAL

AKEEGA

+	JANSSEN BIOTECH	500MG;EQ 50MG BASE	N216793	001	Aug 11, 2023
+	!	500MG;EQ 100MG BASE	N216793	002	Aug 11, 2023

ABROCITINIB

TABLET; ORAL

CIBINQO

+	PFIZER	50MG	N213871	001	Jan 14, 2022
+		100MG	N213871	002	Jan 14, 2022
+	!	200MG	N213871	003	Jan 14, 2022

ACALABRUTINIB

CAPSULE; ORAL

CALQUENCE

+	ASTRAZENECA	100MG	N210259	001	Oct 31, 2017
---	-------------	-------	---------	-----	--------------

ACALABRUTINIB MALEATE

TABLET; ORAL

CALQUENCE

+	ASTRAZENECA	EQ 100MG BASE	N216387	001	Aug 03, 2022
---	-------------	---------------	---------	-----	--------------

ACAMPROSATE CALCIUM

TABLET, DELAYED RELEASE; ORAL

ACAMPROSATE CALCIUM

<u>AB</u>	MYLAN	<u>333MG</u>	<u>A200142</u>	<u>001</u>	Mar 11, 2014
<u>AB</u>	!	ZYDUS PHARMS	<u>333MG</u>	<u>A205995</u>	<u>001</u> May 26, 2017

ACARBOSE

TABLET; ORAL

ACARBOSE

<u>AB</u>	AVET LIFESCIENCES	<u>25MG</u>	<u>A202271</u>	<u>001</u>	Feb 07, 2012
<u>AB</u>		<u>50MG</u>	<u>A202271</u>	<u>002</u>	Feb 07, 2012
<u>AB</u>		<u>100MG</u>	<u>A202271</u>	<u>003</u>	Feb 07, 2012
<u>AB</u>	HIKMA	<u>25MG</u>	<u>A078470</u>	<u>001</u>	May 07, 2008
<u>AB</u>		<u>50MG</u>	<u>A078470</u>	<u>002</u>	May 07, 2008
<u>AB</u>		<u>100MG</u>	<u>A078470</u>	<u>003</u>	May 07, 2008
<u>AB</u>	IMPAX LABS	<u>25MG</u>	<u>A078441</u>	<u>001</u>	May 14, 2009
<u>AB</u>		<u>50MG</u>	<u>A078441</u>	<u>002</u>	May 14, 2009
<u>AB</u>		<u>100MG</u>	<u>A078441</u>	<u>003</u>	May 14, 2009
<u>AB</u>	SOMERSET THERAPS LLC	<u>25MG</u>	<u>A091343</u>	<u>001</u>	Oct 17, 2013
<u>AB</u>		<u>50MG</u>	<u>A091343</u>	<u>002</u>	Oct 17, 2013
<u>AB</u>		<u>100MG</u>	<u>A091343</u>	<u>003</u>	Oct 17, 2013
<u>AB</u>	!	STRIDES PHARMA	<u>25MG</u>	<u>A090912</u>	<u>001</u> Jul 27, 2011
<u>AB</u>		<u>50MG</u>	<u>A090912</u>	<u>002</u>	Jul 27, 2011
<u>AB</u>		<u>100MG</u>	<u>A090912</u>	<u>003</u>	Jul 27, 2011
<u>AB</u>	WATSON LABS	<u>25MG</u>	<u>A077532</u>	<u>001</u>	May 07, 2008
<u>AB</u>		<u>50MG</u>	<u>A077532</u>	<u>002</u>	May 07, 2008
<u>AB</u>		<u>100MG</u>	<u>A077532</u>	<u>003</u>	May 07, 2008

ACEBUTOLOL HYDROCHLORIDE

CAPSULE; ORAL

ACEBUTOLOL HYDROCHLORIDE

<u>AB</u>	!	AMNEAL PHARM	<u>EQ 200MG BASE</u>	<u>A075047</u>	<u>001</u> Dec 30, 1999
<u>AB</u>	!		<u>EQ 400MG BASE</u>	<u>A075047</u>	<u>002</u> Dec 30, 1999
<u>AB</u>		ANI PHARMS	<u>EQ 200MG BASE</u>	<u>A074007</u>	<u>001</u> Oct 18, 1995
<u>AB</u>			<u>EQ 400MG BASE</u>	<u>A074007</u>	<u>002</u> Oct 18, 1995

ACETAMINOPHEN

SOLUTION; INTRAVENOUS

ACETAMINOPHEN

<u>AP</u>	ASPIRO	<u>1GM/100ML (10MG/ML)</u>	<u>A216617</u>	<u>001</u>	Jul 27, 2022
<u>AP</u>	BAXTER HLTHCARE CORP	<u>1GM/100ML (10MG/ML)</u>	<u>A214331</u>	<u>001</u>	Sep 17, 2021
<u>AP</u>	EUGIA PHARMA	<u>1GM/100ML (10MG/ML)</u>	<u>A210969</u>	<u>001</u>	Oct 21, 2020

PRESCRIPTION DRUG PRODUCT LIST

ACETAMINOPHEN

SOLUTION; INTRAVENOUS

ACETAMINOPHEN

<u>AP</u>	HIKMA	<u>1GM/100ML (10MG/ML)</u>	<u>A202605 001</u>	Jun 13, 2016
<u>AP</u>	INFORLIFE	<u>1GM/100ML (10MG/ML)</u>	<u>A215403 001</u>	May 03, 2023
<u>AP</u>	MYLAN	<u>1GM/100ML (10MG/ML)</u>	<u>A213255 001</u>	Aug 07, 2020
<u>AP</u>	! SANDOZ	<u>1GM/100ML (10MG/ML)</u>	<u>A204052 001</u>	Mar 22, 2016
<u>AP</u>	WOCKHARDT BIO AG	<u>1GM/100ML (10MG/ML)</u>	<u>A205746 001</u>	Apr 18, 2023
	+! B BRAUN MEDICAL INC	500MG/50ML (10MG/ML)	N204957 001	Feb 18, 2021
	+!	1GM/100ML (10MG/ML)	N204957 002	Feb 18, 2021
	FRESENIUS KABI USA	1GM/100ML (10MG/ML)	N204767 001	Oct 28, 2015
	+! HIKMA	1GM/100ML (10MG/ML)	N206968 001	Jun 03, 2022

ACETAMINOPHEN; BUTALBITAL

CAPSULE; ORAL

BUTALBITAL AND ACETAMINOPHEN

<u>AA</u>	! DR REDDYS LABS SA	<u>300MG; 50MG</u>	<u>A207313 001</u>	Dec 27, 2017
<u>AA</u>	GRANULES	<u>300MG; 50MG</u>	<u>A213115 001</u>	Nov 22, 2019

TABLET; ORAL

ALLZITAL

<u>AA</u>	! LARKEN LABS INC	<u>325MG; 25MG</u>	<u>A203484 001</u>	Dec 04, 2015
-----------	-------------------	--------------------	--------------------	--------------

BUTALBITAL AND ACETAMINOPHEN

<u>AA</u>	LARKEN LABS INC	<u>325MG; 50MG</u>	<u>A203484 002</u>	Dec 04, 2015
<u>AA</u>	! LGM PHARMA	<u>300MG; 50MG</u>	<u>A090956 001</u>	Aug 23, 2011
<u>AA</u>	MIKART	<u>300MG; 50MG</u>	<u>A207386 001</u>	Nov 15, 2016
<u>AA</u>	NE RX PHARMA	<u>300MG; 50MG</u>	<u>A214955 001</u>	Aug 16, 2022
<u>AA</u>	QUAGEN	<u>300MG; 50MG</u>	<u>A214305 001</u>	Feb 01, 2024
<u>AA</u>		<u>325MG; 50MG</u>	<u>A214291 001</u>	Jan 18, 2024
<u>AA</u>	SENORES PHARMS	<u>300MG; 50MG</u>	<u>A214088 001</u>	Apr 07, 2022
<u>AA</u>		<u>325MG; 25MG</u>	<u>A214088 002</u>	Apr 07, 2022
<u>AA</u>		<u>325MG; 50MG</u>	<u>A214088 003</u>	Apr 07, 2022

BUTAPAP

<u>AA</u>	! MIKART	<u>325MG; 50MG</u>	<u>A089987 001</u>	Oct 26, 1992
-----------	----------	--------------------	--------------------	--------------

ACETAMINOPHEN; BUTALBITAL; CAFFEINE

CAPSULE; ORAL

BUTALBITAL, ACETAMINOPHEN AND CAFFEINE

<u>AA</u>	AUROLIFE PHARMA LLC	<u>325MG; 50MG; 40MG</u>	<u>A204733 001</u>	Sep 26, 2018
<u>AA</u>	DR REDDYS LABS SA	<u>300MG; 50MG; 40MG</u>	<u>A210817 001</u>	Dec 17, 2019
<u>AA</u>	!	<u>325MG; 50MG; 40MG</u>	<u>A089007 001</u>	Mar 17, 1986
<u>AA</u>	GRANULES	<u>300MG; 50MG; 40MG</u>	<u>A213321 001</u>	Apr 08, 2020
<u>AA</u>	LANNETT CO INC	<u>300MG; 50MG; 40MG</u>	<u>A212082 001</u>	Dec 17, 2019
<u>AA</u>		<u>325MG; 50MG; 40MG</u>	<u>A212083 001</u>	Dec 17, 2019
<u>AA</u>	! LGM PHARMA	<u>300MG; 50MG; 40MG</u>	<u>A040885 001</u>	Nov 16, 2009
<u>AA</u>	NOVAST LABS	<u>300MG; 50MG; 40MG</u>	<u>A215047 001</u>	Nov 17, 2021
<u>AA</u>	NUVO PHARMS INC	<u>300MG; 50MG; 40MG</u>	<u>A207118 001</u>	Oct 28, 2016
<u>AA</u>	QUAGEN	<u>300MG; 50MG; 40MG</u>	<u>A214288 001</u>	Feb 08, 2024
<u>AA</u>		<u>325MG; 50MG; 40MG</u>	<u>A216376 001</u>	Mar 12, 2024
<u>AA</u>	SENORES PHARMS	<u>300MG; 50MG; 40MG</u>	<u>A214087 001</u>	Aug 13, 2021
<u>AA</u>		<u>325MG; 50MG; 40MG</u>	<u>A214087 002</u>	Aug 13, 2021
<u>AA</u>	TARO	<u>300MG; 50MG; 40MG</u>	<u>A213046 001</u>	Jul 01, 2020

SOLUTION; ORAL

BUTALBITAL, ACETAMINOPHEN AND CAFFEINE

!	MIKART	325MG/15ML; 50MG/15ML; 40MG/15ML	A040387 001	Jan 31, 2003
---	--------	----------------------------------	-------------	--------------

TABLET; ORAL

BUTALBITAL, ACETAMINOPHEN AND CAFFEINE

<u>AA</u>	ABHAI LLC	<u>325MG; 50MG; 40MG</u>	<u>A211106 001</u>	Sep 26, 2018
<u>AA</u>	+ ACTAVIS LABS UT INC	<u>325MG; 50MG; 40MG</u>	<u>A088616 001</u>	Nov 09, 1984
<u>AA</u>	GRANULES	<u>325MG; 50MG; 40MG</u>	<u>A040864 001</u>	Dec 01, 2008
<u>AA</u>	LANNETT CO INC	<u>325MG; 50MG; 40MG</u>	<u>A200243 001</u>	Sep 13, 2012
<u>AA</u>	LGM PHARMA	<u>325MG; 50MG; 40MG</u>	<u>A209587 001</u>	Oct 31, 2018
<u>AA</u>	MIKART	<u>325MG; 50MG; 40MG</u>	<u>A089175 001</u>	Jan 21, 1987
<u>AA</u>	QUAGEN	<u>325MG; 50MG; 40MG</u>	<u>A214287 001</u>	Jan 18, 2024
<u>AA</u>	! STRIDES PHARMA	<u>325MG; 50MG; 40MG</u>	<u>A040511 001</u>	Aug 27, 2003
<u>AA</u>		<u>325MG; 50MG; 40MG</u>	<u>A203647 001</u>	Sep 21, 2020

ACETAMINOPHEN; BUTALBITAL; CAFFEINE; CODEINE PHOSPHATE

CAPSULE; ORAL

BUTALBITAL, ACETAMINOPHEN, CAFFEINE AND CODEINE PHOSPHATE

<u>AB</u>	HIKMA	<u>300MG; 50MG; 40MG; 30MG</u>	<u>A215138 002</u>	Jan 26, 2022
<u>AB</u>		<u>325MG; 50MG; 40MG; 30MG</u>	<u>A215138 001</u>	Jan 26, 2022
<u>AB</u>	LGM PHARMA	<u>300MG; 50MG; 40MG; 30MG</u>	<u>A076560 002</u>	Jul 19, 2012
<u>AB</u>		<u>325MG; 50MG; 40MG; 30MG</u>	<u>A076560 001</u>	Jun 10, 2004
<u>AB</u>	STRIDES PHARMA	<u>325MG; 50MG; 40MG; 30MG</u>	<u>A204649 001</u>	Jul 08, 2020

PRESCRIPTION DRUG PRODUCT LIST

ACETAMINOPHEN; BUTALBITAL; CAFFEINE; CODEINE PHOSPHATE

CAPSULE;ORAL

FIORICET W/ CODEINE

AB	+ !	ACTAVIS LABS UT INC	325MG;50MG;40MG;30MG	N020232 001	Jul 30, 1992
-----------	------------	---------------------	-----------------------------	--------------------	--------------

ACETAMINOPHEN; CODEINE PHOSPHATE

SOLUTION;ORAL

ACETAMINOPHEN AND CODEINE PHOSPHATE

AA		CHARTWELL	120MG/5ML;12MG/5ML	A089450 001	Oct 27, 1992
-----------	--	-----------	---------------------------	--------------------	--------------

AA	!	GENUS LIFESCIENCES	120MG/5ML;12MG/5ML	A087508 001	
-----------	----------	--------------------	---------------------------	--------------------	--

TABLET;ORAL

ACETAMINOPHEN AND CODEINE PHOSPHATE

AA		AMNEAL PHARMS NY	300MG;30MG	A040779 001	May 29, 2008
-----------	--	------------------	-------------------	--------------------	--------------

AA		AUROLIFE PHARMA LLC	300MG;15MG	A202800 001	Apr 15, 2013
-----------	--	---------------------	-------------------	--------------------	--------------

AA			300MG;30MG	A202800 002	Apr 15, 2013
-----------	--	--	-------------------	--------------------	--------------

AA			300MG;60MG	A202800 003	Apr 15, 2013
-----------	--	--	-------------------	--------------------	--------------

AA		ELITE LABS INC	300MG;15MG	A212418 001	Sep 10, 2019
-----------	--	----------------	-------------------	--------------------	--------------

AA			300MG;30MG	A212418 002	Sep 10, 2019
-----------	--	--	-------------------	--------------------	--------------

AA			300MG;60MG	A212418 003	Sep 10, 2019
-----------	--	--	-------------------	--------------------	--------------

AA	!	SPECGX LLC	300MG;15MG	A040419 001	May 31, 2001
-----------	----------	------------	-------------------	--------------------	--------------

AA	!		300MG;30MG	A040419 002	May 31, 2001
-----------	----------	--	-------------------	--------------------	--------------

AA	!		300MG;60MG	A040419 003	May 31, 2001
-----------	----------	--	-------------------	--------------------	--------------

AA		STRIDES PHARMA	300MG;15MG	A089990 001	Sep 30, 1988
-----------	--	----------------	-------------------	--------------------	--------------

AA			300MG;30MG	A089805 001	Sep 30, 1988
-----------	--	--	-------------------	--------------------	--------------

AA			300MG;60MG	A089828 001	Sep 30, 1988
-----------	--	--	-------------------	--------------------	--------------

AA		SUN PHARM INDS LTD	300MG;60MG	A087083 001	
-----------	--	--------------------	-------------------	--------------------	--

AA		WES PHARMA INC	300MG;15MG	A211610 001	Jun 27, 2019
-----------	--	----------------	-------------------	--------------------	--------------

AA			300MG;30MG	A211610 002	Jun 27, 2019
-----------	--	--	-------------------	--------------------	--------------

AA			300MG;60MG	A211610 003	Jun 27, 2019
-----------	--	--	-------------------	--------------------	--------------

ACETAMINOPHEN; HYDROCODONE BITARTRATE

SOLUTION;ORAL

HYDROCODONE BITARTRATE AND ACETAMINOPHEN

AA	!	CHARTWELL	325MG/15ML;7.5MG/15ML	A040482 001	Sep 25, 2003
-----------	----------	-----------	------------------------------	--------------------	--------------

AA		GENUS	325MG/15ML;7.5MG/15ML	A040894 001	Jul 19, 2011
-----------	--	-------	------------------------------	--------------------	--------------

AA		PHARM ASSOC	325MG/15ML;7.5MG/15ML	A040838 001	May 10, 2013
-----------	--	-------------	------------------------------	--------------------	--------------

AA		WES PHARMA INC	325MG/15ML;7.5MG/15ML	A211023 001	Mar 08, 2019
-----------	--	----------------	------------------------------	--------------------	--------------

AA	!	MIKART	300MG/15ML;10MG/15ML	A040881 001	Feb 25, 2010
-----------	----------	--------	-----------------------------	--------------------	--------------

AA	!	PHARM ASSOC	325MG/15ML;10MG/15ML	A040834 001	Apr 18, 2008
-----------	----------	-------------	-----------------------------	--------------------	--------------

TABLET;ORAL

ANEXSIA 5/325

AA	!	SPECGX LLC	325MG;5MG	A040409 001	Oct 20, 2000
-----------	----------	------------	------------------	--------------------	--------------

ANEXSIA 7.5/325

AA	!	SPECGX LLC	325MG;7.5MG	A040405 001	Sep 08, 2000
-----------	----------	------------	--------------------	--------------------	--------------

HYDROCODONE BITARTRATE AND ACETAMINOPHEN

AA		ABHAI LLC	300MG;5MG	A209036 001	Jun 21, 2017
-----------	--	-----------	------------------	--------------------	--------------

AA			300MG;7.5MG	A209036 002	Jun 21, 2017
-----------	--	--	--------------------	--------------------	--------------

AA			300MG;10MG	A209036 003	Jun 21, 2017
-----------	--	--	-------------------	--------------------	--------------

AA			325MG;5MG	A209037 001	Jun 21, 2017
-----------	--	--	------------------	--------------------	--------------

AA			325MG;7.5MG	A209037 002	Jun 21, 2017
-----------	--	--	--------------------	--------------------	--------------

AA			325MG;10MG	A209037 003	Jun 21, 2017
-----------	--	--	-------------------	--------------------	--------------

AA		AMNEAL PHARMS NY	325MG;5MG	A040736 001	Aug 25, 2006
-----------	--	------------------	------------------	--------------------	--------------

AA			325MG;7.5MG	A040746 002	May 10, 2016
-----------	--	--	--------------------	--------------------	--------------

AA			325MG;10MG	A040746 001	Aug 25, 2006
-----------	--	--	-------------------	--------------------	--------------

AA	!	ASCENT PHARMS INC	325MG;2.5MG	A211487 001	Nov 07, 2018
-----------	----------	-------------------	--------------------	--------------------	--------------

AA			325MG;5MG	A211487 002	Nov 07, 2018
-----------	--	--	------------------	--------------------	--------------

AA			325MG;7.5MG	A211487 003	Nov 07, 2018
-----------	--	--	--------------------	--------------------	--------------

AA			325MG;10MG	A211487 004	Nov 07, 2018
-----------	--	--	-------------------	--------------------	--------------

AA		AUROLIFE PHARMA LLC	300MG;5MG	A207709 001	Sep 13, 2018
-----------	--	---------------------	------------------	--------------------	--------------

AA			300MG;7.5MG	A207709 002	Sep 13, 2018
-----------	--	--	--------------------	--------------------	--------------

AA			300MG;10MG	A207709 003	Sep 13, 2018
-----------	--	--	-------------------	--------------------	--------------

AA			325MG;5MG	A201013 001	Apr 11, 2012
-----------	--	--	------------------	--------------------	--------------

AA			325MG;7.5MG	A201013 002	Apr 11, 2012
-----------	--	--	--------------------	--------------------	--------------

AA			325MG;10MG	A201013 003	Apr 11, 2012
-----------	--	--	-------------------	--------------------	--------------

AA	!	CHARTWELL	300MG;5MG	A040658 001	Jan 19, 2006
-----------	----------	-----------	------------------	--------------------	--------------

AA	!		300MG;7.5MG	A040658 002	Mar 24, 2006
-----------	----------	--	--------------------	--------------------	--------------

AA	!		300MG;10MG	A040658 003	Jun 23, 2004
-----------	----------	--	-------------------	--------------------	--------------

AA			325MG;7.5MG	A040432 001	Jan 22, 2003
-----------	--	--	--------------------	--------------------	--------------

AA		ELITE LABS INC	325MG;2.5MG	A209924 001	Nov 16, 2018
-----------	--	----------------	--------------------	--------------------	--------------

AA			325MG;5MG	A209924 002	Nov 16, 2018
-----------	--	--	------------------	--------------------	--------------

AA			325MG;7.5MG	A209924 003	Nov 16, 2018
-----------	--	--	--------------------	--------------------	--------------

AA			325MG;10MG	A209924 004	Nov 16, 2018
-----------	--	--	-------------------	--------------------	--------------

AA		EPIC PHARMA LLC	325MG;5MG	A203863 001	Mar 30, 2018
-----------	--	-----------------	------------------	--------------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

ACETAMINOPHEN; HYDROCODONE BITARTRATE

TABLET; ORAL

HYDROCODONE BITARTRATE AND ACETAMINOPHEN

<u>AA</u>		<u>325MG; 7.5MG</u>	<u>A203863 002</u>	Mar 30, 2018
<u>AA</u>		<u>325MG; 10MG</u>	<u>A203863 003</u>	Mar 30, 2018
<u>AA</u>	GRANULES	<u>325MG; 5MG</u>	<u>A211729 001</u>	Jan 03, 2020
<u>AA</u>		<u>325MG; 7.5MG</u>	<u>A211729 002</u>	Jan 03, 2020
<u>AA</u>		<u>325MG; 10MG</u>	<u>A211729 003</u>	Jan 03, 2020
<u>AA</u>	RHODES PHARMS	<u>300MG; 5MG</u>	<u>A207808 001</u>	Mar 30, 2018
<u>AA</u>		<u>300MG; 7.5MG</u>	<u>A207808 002</u>	Mar 30, 2018
<u>AA</u>		<u>300MG; 10MG</u>	<u>A207808 003</u>	Mar 30, 2018
<u>AA</u>		<u>325MG; 5MG</u>	<u>A202991 001</u>	Apr 12, 2016
<u>AA</u>		<u>325MG; 7.5MG</u>	<u>A202991 002</u>	Apr 12, 2016
<u>AA</u>		<u>325MG; 10MG</u>	<u>A202991 003</u>	Apr 12, 2016
<u>AA</u>	SPECGX LLC	<u>300MG; 5MG</u>	<u>A206718 001</u>	Mar 31, 2017
<u>AA</u>		<u>300MG; 7.5MG</u>	<u>A206718 002</u>	Mar 31, 2017
<u>AA</u>		<u>300MG; 10MG</u>	<u>A206718 003</u>	Mar 31, 2017
<u>AA</u>	!	<u>325MG; 10MG</u>	<u>A040400 001</u>	Jul 26, 2000
<u>AA</u>	STRIDES PHARMA	<u>325MG; 5MG</u>	<u>A040655 001</u>	Jan 19, 2006
<u>AA</u>		<u>325MG; 5MG</u>	<u>A202935 002</u>	Jun 15, 2016
<u>AA</u>		<u>325MG; 7.5MG</u>	<u>A040656 001</u>	Jan 19, 2006
<u>AA</u>		<u>325MG; 7.5MG</u>	<u>A202935 003</u>	Jun 15, 2016
<u>AA</u>		<u>325MG; 10MG</u>	<u>A040355 001</u>	May 31, 2000
<u>AA</u>		<u>325MG; 10MG</u>	<u>A202935 004</u>	Jun 15, 2016
<u>AA</u>	TRIS PHARMA INC	<u>300MG; 5MG</u>	<u>A202214 004</u>	Mar 15, 2016
<u>AA</u>		<u>300MG; 7.5MG</u>	<u>A202214 005</u>	Mar 15, 2016
<u>AA</u>		<u>300MG; 10MG</u>	<u>A202214 006</u>	Mar 15, 2016
<u>AA</u>		<u>325MG; 5MG</u>	<u>A202214 001</u>	Mar 27, 2013
<u>AA</u>		<u>325MG; 7.5MG</u>	<u>A202214 002</u>	Mar 27, 2013
<u>AA</u>		<u>325MG; 10MG</u>	<u>A202214 003</u>	Mar 27, 2013
<u>AA</u>	WES PHARMA INC	<u>300MG; 5MG</u>	<u>A207509 001</u>	Oct 29, 2018
<u>AA</u>		<u>300MG; 7.5MG</u>	<u>A207509 002</u>	Oct 29, 2018
<u>AA</u>		<u>300MG; 10MG</u>	<u>A207509 003</u>	Oct 29, 2018
<u>AA</u>		<u>325MG; 5MG</u>	<u>A210211 001</u>	Oct 30, 2017
<u>AA</u>		<u>325MG; 7.5MG</u>	<u>A210211 002</u>	Oct 30, 2017
<u>AA</u>		<u>325MG; 10MG</u>	<u>A210211 003</u>	Oct 30, 2017

ACETAMINOPHEN; IBUPROFEN

TABLET; ORAL

COMBOGESIC

+	!	AFT PHARMS US	325MG; 97.5MG	N209471 001	Mar 01, 2023
---	---	---------------	---------------	-------------	--------------

ACETAMINOPHEN; IBUPROFEN SODIUM

SOLUTION; INTRAVENOUS

COMBOGESIC IV

+	!	HIKMA	1GM/100ML (10MG/ML); EQ 300MG BASE/100ML (EQ 3MG BASE/ML)	N215320 001	Oct 17, 2023
---	---	-------	---	-------------	--------------

ACETAMINOPHEN; OXYCODONE HYDROCHLORIDE

SOLUTION; ORAL

OXYCODONE AND ACETAMINOPHEN

<u>AA</u>	!	ABHAI LLC	<u>325MG/5ML; 5MG/5ML</u>	<u>A211499 001</u>	Dec 31, 2018
<u>AA</u>		NOSTRUM LABS INC	<u>325MG/5ML; 5MG/5ML</u>	<u>A201448 001</u>	Aug 26, 2021
		MIKART	300MG/5ML; 10MG/5ML	A202142 001	Nov 27, 2018

TABLET; ORAL

OXYCET

<u>AA</u>		SPECGX LLC	<u>325MG; 5MG</u>	<u>A087463 001</u>	Dec 07, 1983
-----------	--	------------	-------------------	--------------------	--------------

OXYCODONE AND ACETAMINOPHEN

<u>AA</u>		ABHAI LLC	<u>325MG; 2.5MG</u>	<u>A210644 001</u>	Feb 09, 2018
<u>AA</u>			<u>325MG; 5MG</u>	<u>A210644 002</u>	Feb 09, 2018
<u>AA</u>			<u>325MG; 7.5MG</u>	<u>A210644 003</u>	Feb 09, 2018
<u>AA</u>			<u>325MG; 10MG</u>	<u>A210644 004</u>	Feb 09, 2018
<u>AA</u>	ACTAVIS ELIZABETH		<u>325MG; 2.5MG</u>	<u>A201447 001</u>	Apr 12, 2013
<u>AA</u>			<u>325MG; 5MG</u>	<u>A201447 002</u>	Apr 12, 2013
<u>AA</u>			<u>325MG; 7.5MG</u>	<u>A201447 003</u>	Apr 12, 2013
<u>AA</u>			<u>325MG; 10MG</u>	<u>A201447 004</u>	Apr 12, 2013
<u>AA</u>	ALVOGEN		<u>325MG; 5MG</u>	<u>A202677 003</u>	Mar 08, 2016
<u>AA</u>			<u>325MG; 7.5MG</u>	<u>A202677 001</u>	Jul 26, 2012
<u>AA</u>			<u>325MG; 10MG</u>	<u>A202677 002</u>	Jul 26, 2012
<u>AA</u>	AMNEAL PHARMS		<u>325MG; 5MG</u>	<u>A040777 001</u>	Nov 27, 2007
<u>AA</u>	AMNEAL PHARMS NY		<u>325MG; 7.5MG</u>	<u>A040778 002</u>	Jun 27, 2014
<u>AA</u>			<u>325MG; 10MG</u>	<u>A040778 001</u>	Nov 27, 2007
<u>AA</u>	ASCENT PHARMS INC		<u>325MG; 2.5MG</u>	<u>A207419 001</u>	Mar 22, 2017
<u>AA</u>			<u>325MG; 5MG</u>	<u>A207419 002</u>	Mar 22, 2017

PRESCRIPTION DRUG PRODUCT LIST

ACETAMINOPHEN; OXYCODONE HYDROCHLORIDE

TABLET; ORAL

OXYCODONE AND ACETAMINOPHEN

<u>AA</u>		<u>325MG; 7.5MG</u>	<u>A207419 003</u>	Mar 22, 2017
<u>AA</u>		<u>325MG; 10MG</u>	<u>A207419 004</u>	Mar 22, 2017
<u>AA</u>	AUROLIFE PHARMA LLC	<u>325MG; 2.5MG</u>	<u>A201972 001</u>	Jul 15, 2013
<u>AA</u>		<u>325MG; 5MG</u>	<u>A201972 002</u>	Jul 15, 2013
<u>AA</u>		<u>325MG; 7.5MG</u>	<u>A201972 003</u>	Jul 15, 2013
<u>AA</u>		<u>325MG; 10MG</u>	<u>A201972 004</u>	Jul 15, 2013
<u>AA</u>	CHARTWELL	<u>325MG; 5MG</u>	<u>A207834 001</u>	Aug 15, 2019
<u>AA</u>		<u>325MG; 7.5MG</u>	<u>A207834 002</u>	Aug 15, 2019
<u>AA</u>		<u>325MG; 10MG</u>	<u>A207834 003</u>	Aug 15, 2019
<u>AA</u>	ELITE LABS INC	<u>325MG; 5MG</u>	<u>A209385 001</u>	Jul 02, 2018
<u>AA</u>		<u>325MG; 7.5MG</u>	<u>A209385 002</u>	Jul 02, 2018
<u>AA</u>		<u>325MG; 10MG</u>	<u>A209385 003</u>	Jul 02, 2018
<u>AA</u>	EPIC PHARMA LLC	<u>325MG; 5MG</u>	<u>A203864 001</u>	Jul 02, 2018
<u>AA</u>		<u>325MG; 7.5MG</u>	<u>A203864 002</u>	Jul 02, 2018
<u>AA</u>		<u>325MG; 10MG</u>	<u>A203864 003</u>	Jul 02, 2018
<u>AA</u>	GRANULES	<u>325MG; 2.5MG</u>	<u>A211708 001</u>	Oct 31, 2019
<u>AA</u>		<u>325MG; 5MG</u>	<u>A211708 002</u>	Oct 31, 2019
<u>AA</u>		<u>325MG; 7.5MG</u>	<u>A211708 003</u>	Oct 31, 2019
<u>AA</u>		<u>325MG; 10MG</u>	<u>A211708 004</u>	Oct 31, 2019
<u>AA</u>	NOVEL LABS INC	<u>325MG; 2.5MG</u>	<u>A204407 001</u>	Feb 24, 2017
<u>AA</u>		<u>325MG; 5MG</u>	<u>A204407 002</u>	Feb 24, 2017
<u>AA</u>		<u>325MG; 7.5MG</u>	<u>A204407 003</u>	Feb 24, 2017
<u>AA</u>		<u>325MG; 10MG</u>	<u>A204407 004</u>	Feb 24, 2017
<u>AA</u>	RHODES PHARMS	<u>325MG; 5MG</u>	<u>A201278 001</u>	Aug 28, 2014
<u>AA</u>		<u>325MG; 7.5MG</u>	<u>A201278 002</u>	Aug 28, 2014
<u>AA</u>		<u>325MG; 10MG</u>	<u>A201278 003</u>	Aug 28, 2014
<u>AA</u>	SPECGX LLC	<u>325MG; 7.5MG</u>	<u>A040545 001</u>	Jun 30, 2004
<u>AA</u>		<u>325MG; 10MG</u>	<u>A040545 002</u>	Jun 30, 2004
<u>AA</u>	WES PHARMA INC	<u>325MG; 5MG</u>	<u>A207510 001</u>	Mar 21, 2018
<u>AA</u>		<u>325MG; 7.5MG</u>	<u>A207510 002</u>	Mar 21, 2018
<u>AA</u>		<u>325MG; 10MG</u>	<u>A207510 003</u>	Mar 21, 2018

PERCOCET

<u>AA</u>	!	ENDO OPERATIONS	<u>325MG; 2.5MG</u>	<u>A040330 001</u>	Jun 25, 1999
<u>AA</u>	!		<u>325MG; 5MG</u>	<u>A040330 002</u>	Jun 25, 1999
<u>AA</u>	!		<u>325MG; 7.5MG</u>	<u>A040330 003</u>	Nov 23, 2001
<u>AA</u>	!		<u>325MG; 10MG</u>	<u>A040330 004</u>	Nov 23, 2001

OXYCODONE AND ACETAMINOPHEN

!	MIKART	300MG; 2.5MG	A040608 001	Dec 30, 2005
!		300MG; 5MG	A040608 002	Dec 30, 2005
!		300MG; 7.5MG	A040608 003	Dec 30, 2005
!		300MG; 10MG	A040608 004	Dec 30, 2005

ACETAMINOPHEN; TRAMADOL HYDROCHLORIDE

TABLET; ORAL

TRAMADOL HYDROCHLORIDE AND ACETAMINOPHEN

<u>AB</u>	ALKEM LABS LTD	<u>325MG; 37.5MG</u>	<u>A202076 001</u>	Mar 30, 2012
<u>AB</u>	!	AMNEAL PHARMS	<u>A090485 001</u>	Dec 09, 2009
<u>AB</u>	AUROBINDO PHARMA	<u>325MG; 37.5MG</u>	<u>A207152 001</u>	Mar 22, 2017
<u>AB</u>	MICRO LABS LTD	<u>325MG; 37.5MG</u>	<u>A201952 001</u>	Dec 14, 2012
	INDIA			
<u>AB</u>	RISING	<u>325MG; 37.5MG</u>	<u>A077858 001</u>	Sep 26, 2008
<u>AB</u>	ZYDUS PHARMS USA	<u>325MG; 37.5MG</u>	<u>A090460 001</u>	Sep 06, 2012
	INC			

ACETAZOLAMIDE

CAPSULE, EXTENDED RELEASE; ORAL

ACETAZOLAMIDE

<u>AB</u>	ACCORD HLTHCARE	<u>500MG</u>	<u>A207659 001</u>	Oct 18, 2018
<u>AB</u>	ALEMBIC	<u>500MG</u>	<u>A210423 001</u>	Feb 19, 2019
<u>AB</u>	!	CADILA	<u>A205301 001</u>	Jan 16, 2019
<u>AB</u>	HERITAGE PHARMA	<u>500MG</u>	<u>A040904 001</u>	Dec 10, 2008
<u>AB</u>	INDICUS PHARMA	<u>500MG</u>	<u>A090779 001</u>	Jul 14, 2011
<u>AB</u>	MICRO LABS LTD	<u>500MG</u>	<u>A207401 001</u>	Oct 01, 2020
	INDIA			
<u>AB</u>	NOSTRUM LABS INC	<u>500MG</u>	<u>A204691 001</u>	Mar 29, 2016
<u>AB</u>	NOVAST LABS	<u>500MG</u>	<u>A203434 001</u>	Sep 30, 2016

TABLET; ORAL

ACETAZOLAMIDE

<u>AB</u>	APPCO	<u>125MG</u>	<u>A211372 001</u>	Feb 22, 2021
<u>AB</u>		<u>250MG</u>	<u>A211372 002</u>	Feb 22, 2021
<u>AB</u>	CHARTWELL MOLECULAR	<u>250MG</u>	<u>A084840 001</u>	

PRESCRIPTION DRUG PRODUCT LIST

ACETAZOLAMIDE

TABLET; ORAL

ACETAZOLAMIDE

<u>AB</u>	EPIC PHARMA LLC	<u>125MG</u>	<u>A218023 001</u>	May 14, 2024
<u>AB</u>		<u>250MG</u>	<u>A218023 002</u>	May 14, 2024
<u>AB</u>	HERITAGE PHARMA	<u>125MG</u>	<u>A205530 001</u>	Oct 27, 2016
<u>AB</u>		<u>250MG</u>	<u>A205530 002</u>	Oct 27, 2016
<u>AB</u>	MANKIND PHARMA	<u>125MG</u>	<u>A214282 001</u>	Oct 07, 2020
<u>AB</u>		<u>250MG</u>	<u>A214282 002</u>	Oct 07, 2020
<u>AB</u>	NE RX PHARMA	<u>125MG</u>	<u>A217197 001</u>	May 09, 2023
<u>AB</u>		<u>250MG</u>	<u>A217197 002</u>	May 09, 2023
<u>AB</u>	NOVITIUM PHARMA	<u>125MG</u>	<u>A210588 001</u>	Oct 17, 2019
<u>AB</u>		<u>250MG</u>	<u>A210588 002</u>	Oct 17, 2019
<u>AB</u>	RUBICON	<u>125MG</u>	<u>A215101 001</u>	Aug 19, 2021
<u>AB</u>		<u>250MG</u>	<u>A215101 002</u>	Aug 19, 2021
<u>AB</u>	STRIDES PHARMA	<u>125MG</u>	<u>A209734 001</u>	Nov 20, 2017
<u>AB</u>		<u>250MG</u>	<u>A209734 002</u>	Nov 20, 2017
<u>AB</u>	TARO	<u>125MG</u>	<u>A040195 001</u>	May 28, 1997
<u>AB</u>	!	<u>250MG</u>	<u>A040195 002</u>	May 28, 1997
<u>AB</u>	ZYDUS LIFESCIENCES	<u>125MG</u>	<u>A211069 001</u>	Apr 04, 2023
<u>AB</u>		<u>250MG</u>	<u>A211069 002</u>	Apr 04, 2023
BX	HIBROW HLTHCARE	125MG	A211556 001	Oct 18, 2019
BX		250MG	A211556 002	Oct 18, 2019

ACETAZOLAMIDE SODIUM

INJECTABLE; INJECTION

ACETAZOLAMIDE SODIUM

<u>AP</u>	AVET LIFESCIENCES	<u>EQ 500MG BASE/VIAL</u>	<u>A202693 001</u>	Dec 19, 2014
<u>AP</u>	HIKMA	<u>EQ 500MG BASE/VIAL</u>	<u>A040089 001</u>	Feb 28, 1995
<u>AP</u>	MYLAN ASI	<u>EQ 500MG BASE/VIAL</u>	<u>A200880 001</u>	May 09, 2012
<u>AP</u>	!	<u>EQ 500MG BASE/VIAL</u>	<u>A040784 001</u>	Dec 10, 2008
<u>AP</u>	ZYDUS PHARMS	<u>EQ 500MG BASE/VIAL</u>	<u>A206533 001</u>	Apr 15, 2019

ACETIC ACID, GLACIAL

SOLUTION; IRRIGATION, URETHRAL

ACETIC ACID 0.25% IN PLASTIC CONTAINER

<u>AT</u>	B BRAUN	<u>250MG/100ML</u>	<u>N018161 001</u>	
<u>AT</u>	BAXTER HLTHCARE	<u>250MG/100ML</u>	<u>N018523 001</u>	Feb 19, 1982
<u>AT</u>	! ICU MEDICAL INC	<u>250MG/100ML</u>	<u>N017656 001</u>	

SOLUTION/DROPS; OTIC

ACETIC ACID

<u>AT</u>	RISING	<u>2%</u>	<u>A207280 001</u>	Mar 09, 2018
<u>AT</u>	!	<u>2%</u>	<u>A040607 001</u>	Feb 24, 2005
<u>AT</u>	TARO	<u>2%</u>	<u>A088638 001</u>	Sep 06, 1984

ACETIC ACID, GLACIAL; HYDROCORTISONE

SOLUTION/DROPS; OTIC

HYDROCORTISONE AND ACETIC ACID

<u>AT</u>	COSETTE	<u>2%:1%</u>	<u>A040609 001</u>	Feb 06, 2006
<u>AT</u>	! TARO	<u>2%:1%</u>	<u>A088759 001</u>	Mar 04, 1985
	<u>VOSOL HC</u>			
<u>AT</u>	! SAPTALIS PHARMS	<u>2%:1%</u>	<u>N012770 001</u>	

ACETOHYDROXAMIC ACID

TABLET; ORAL

LITHOSTAT

+	MISSION PHARMA	250MG	N018749 001	May 31, 1983
---	----------------	-------	-------------	--------------

ACETYLCHOLINE CHLORIDE

FOR SOLUTION; OPHTHALMIC

MIOCHOL-E

+	BAUSCH AND LOMB	20MG/VIAL	N020213 001	Sep 22, 1993
---	-----------------	-----------	-------------	--------------

ACETYLCYSTEINE

INJECTABLE; INTRAVENOUS

ACETADOTE

<u>AP</u>	! CUMBERLAND PHARMS	<u>6GM/30ML (200MG/ML)</u>	<u>N021539 001</u>	Jan 23, 2004
-----------	---------------------	----------------------------	--------------------	--------------

ACETYLCYSTEINE

<u>AP</u>	EUGIA PHARMA	<u>6GM/30ML (200MG/ML)</u>	<u>A207358 001</u>	Feb 29, 2016
<u>AP</u>	FRESENIUS KABI USA	<u>6GM/30ML (200MG/ML)</u>	<u>A200644 001</u>	Nov 07, 2012
<u>AP</u>	GLENMARK PHARMS INC	<u>6GM/30ML (200MG/ML)</u>	<u>A213693 001</u>	Feb 03, 2022
<u>AP</u>	INDOCO	<u>6GM/30ML (200MG/ML)</u>	<u>A215620 001</u>	Feb 23, 2022
<u>AP</u>	RISING	<u>6GM/30ML (200MG/ML)</u>	<u>A203173 001</u>	Mar 24, 2015
<u>AP</u>	SAGENT PHARMS INC	<u>6GM/30ML (200MG/ML)</u>	<u>A091684 001</u>	Oct 31, 2017
<u>AP</u>	SOMERSET THERAPS	<u>6GM/30ML (200MG/ML)</u>	<u>A218397 001</u>	Jun 07, 2024

PRESCRIPTION DRUG PRODUCT LIST

ACETYLCYSTEINE

INJECTABLE; INTRAVENOUS

ACETYLCYSTEINE

<u>AP</u>	LLC STERISCIENCE	<u>6GM/30ML (200MG/ML)</u>	<u>A217182 001</u>	Apr 19, 2023
<u>AP</u>	SPECLTS ZYDUS PHARMS	<u>6GM/30ML (200MG/ML)</u>	<u>A208166 001</u>	Jul 20, 2018

SOLUTION; INHALATION, ORAL

ACETYLCYSTEINE

<u>AN</u>	ALVOGEN	<u>10%</u>	<u>A204674 001</u>	Feb 11, 2014
<u>AN</u>		<u>20%</u>	<u>A203853 001</u>	Jun 21, 2012
<u>AN</u>	! AM REGENT	<u>10%</u>	<u>A072489 001</u>	Jul 28, 1995
<u>AN</u>	!	<u>20%</u>	<u>A072547 001</u>	Jul 28, 1995
<u>AN</u>	HOSPIRA	<u>10%</u>	<u>A073664 001</u>	Aug 30, 1994
<u>AN</u>		<u>20%</u>	<u>A074037 001</u>	Aug 30, 1994

ACITRETIN

CAPSULE; ORAL

ACITRETIN

<u>AB</u>	ALEMBIC	<u>10MG</u>	<u>A217774 001</u>	Aug 05, 2024
<u>AB</u>		<u>17.5MG</u>	<u>A217774 002</u>	Aug 05, 2024
<u>AB</u>		<u>25MG</u>	<u>A217774 003</u>	Aug 05, 2024
<u>AB</u>	BARR LABS INC	<u>10MG</u>	<u>A091455 001</u>	Apr 04, 2013
<u>AB</u>		<u>25MG</u>	<u>A091455 002</u>	Apr 04, 2013
<u>AB</u>	IMPAX LABS INC	<u>10MG</u>	<u>A202552 001</u>	Dec 23, 2015
<u>AB</u>		<u>17.5MG</u>	<u>A202552 002</u>	Dec 23, 2015
<u>AB</u>		<u>22.5MG</u>	<u>A202552 003</u>	Dec 23, 2015
<u>AB</u>		<u>25MG</u>	<u>A202552 004</u>	Dec 23, 2015
<u>AB</u>	MYLAN	<u>10MG</u>	<u>A202148 001</u>	Sep 10, 2015
<u>AB</u>		<u>25MG</u>	<u>A202148 002</u>	Sep 10, 2015
<u>AB</u>	SIGMAPHARM LABS LLC	<u>10MG</u>	<u>A204633 001</u>	May 22, 2015
<u>AB</u>		<u>17.5MG</u>	<u>A204633 002</u>	May 22, 2015
<u>AB</u>		<u>22.5MG</u>	<u>A204633 003</u>	May 22, 2015
<u>AB</u>	!	<u>25MG</u>	<u>A204633 004</u>	May 22, 2015
<u>AB</u>	TEVA PHARMS USA	<u>17.5MG</u>	<u>A202897 001</u>	Apr 04, 2013
<u>AB</u>		<u>22.5MG</u>	<u>A202897 002</u>	Apr 04, 2013

ACLIDINIUM BROMIDE

POWDER, METERED; INHALATION

TUDORZA PRESSAIR

+	!	COVIS	0.4MG/INH	N202450 001	Jul 23, 2012
---	---	-------	-----------	-------------	--------------

ACLIDINIUM BROMIDE; FORMOTEROL FUMARATE

POWDER, METERED; INHALATION

DUAKLIR PRESSAIR

+	!	COVIS	0.4MG/INH; 0.012MG/INH	N210595 001	Mar 29, 2019
---	---	-------	------------------------	-------------	--------------

ACORAMIDIS HYDROCHLORIDE

TABLET; ORAL

ATTRUBY

+	!	BRIDGEBIO PHARMA	EQ 356MG BASE	N216540 001	Nov 22, 2024
---	---	------------------	---------------	-------------	--------------

ACYCLOVIR

CAPSULE; ORAL

ACYCLOVIR

<u>AB</u>	ADAPTIS	<u>200MG</u>	<u>A075090 001</u>	Jan 26, 1999
<u>AB</u>	! APOTEX	<u>200MG</u>	<u>A075677 001</u>	Sep 28, 2005
<u>AB</u>	CADILA	<u>200MG</u>	<u>A204313 001</u>	Mar 25, 2016
<u>AB</u>	CADILA PHARMS LTD	<u>200MG</u>	<u>A201445 001</u>	Mar 06, 2014
<u>AB</u>	CARLSBAD TECHNOLOGY	<u>200MG</u>	<u>A206261 001</u>	Aug 16, 2017
<u>AB</u>	HERITAGE	<u>200MG</u>	<u>A074889 001</u>	Oct 31, 1997
<u>AB</u>	TEVA	<u>200MG</u>	<u>A074578 001</u>	Apr 22, 1997
<u>AB</u>	YILING	<u>200MG</u>	<u>A212173 001</u>	Sep 14, 2020

CREAM; TOPICAL

ACYCLOVIR

<u>AB</u>	ALEMBIC	<u>5%</u>	<u>A212361 001</u>	Nov 21, 2023
<u>AB</u>	AMNEAL	<u>5%</u>	<u>A208766 001</u>	Nov 09, 2020
<u>AB</u>	! PADAGIS ISRAEL	<u>5%</u>	<u>A208702 001</u>	Feb 04, 2019
<u>AB</u>	TARO	<u>5%</u>	<u>A205470 001</u>	Jun 12, 2024
<u>AB</u>	ZYDUS LIFESCIENCES	<u>5%</u>	<u>A206770 001</u>	Feb 28, 2023

ZOVIRAX

<u>AB</u>	+ BAUSCH	<u>5%</u>	<u>N021478 001</u>	Dec 30, 2002
-----------	----------	-----------	--------------------	--------------

OINTMENT; TOPICAL

ACYCLOVIR

<u>AB</u>	ALEMBIC	<u>5%</u>	<u>A209000 001</u>	Apr 06, 2018
-----------	---------	-----------	--------------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

ACYCLOVIR

OINTMENT; TOPICAL

ACYCLOVIR

<u>AB</u>	AMNEAL PHARMS	<u>5%</u>	<u>A204605</u>	<u>001</u>	Jun 18, 2014
<u>AB</u>	APOTEX	<u>5%</u>	<u>A210774</u>	<u>001</u>	Sep 06, 2019
<u>AB</u>	CHARTWELL RX	<u>5%</u>	<u>A212495</u>	<u>001</u>	Apr 07, 2020
<u>AB</u>	CIPLA	<u>5%</u>	<u>A211794</u>	<u>001</u>	Jan 18, 2019
<u>AB</u>	FOUGERA PHARMS INC	<u>5%</u>	<u>A206633</u>	<u>001</u>	May 11, 2016
<u>AB</u>	GLENMARK SPECLT	<u>5%</u>	<u>A205510</u>	<u>001</u>	Jul 31, 2017
<u>AB</u>	MACLEODS PHARMS LTD	<u>5%</u>	<u>A212444</u>	<u>001</u>	May 19, 2021
<u>AB</u>	MYLAN PHARMS INC	<u>5%</u>	<u>A202459</u>	<u>001</u>	Apr 03, 2013
<u>AB</u>	TARO	<u>5%</u>	<u>A205469</u>	<u>001</u>	Dec 21, 2016
<u>AB</u>	TORRENT	<u>5%</u>	<u>A209971</u>	<u>001</u>	Jan 11, 2019
<u>AB</u>	XIROMED	<u>5%</u>	<u>A201501</u>	<u>001</u>	Jan 29, 2020
<u>AB</u>	ZYDUS LIFESCIENCES	<u>5%</u>	<u>A205974</u>	<u>001</u>	Mar 15, 2019

ZOVIRAX

<u>AB</u>	<u>+</u> ! BAUSCH	<u>5%</u>	<u>N018604</u>	<u>001</u>	Mar 29, 1982
-----------	-------------------	-----------	----------------	------------	--------------

SUSPENSION; ORAL

ACYCLOVIR

<u>AB</u>	ACTAVIS MID ATLANTIC	<u>200MG/5ML</u>	<u>A074738</u>	<u>001</u>	Apr 28, 1997
<u>AB</u>	AUROBINDO PHARMA	<u>200MG/5ML</u>	<u>A216331</u>	<u>001</u>	Mar 31, 2022
<u>AB</u>	CHARTWELL RX	<u>200MG/5ML</u>	<u>A212718</u>	<u>001</u>	Apr 23, 2020
<u>AB</u>	HETERO LABS LTD III	<u>200MG/5ML</u>	<u>A215669</u>	<u>001</u>	Jul 01, 2022
<u>AB</u>	MSN	<u>200MG/5ML</u>	<u>A217393</u>	<u>001</u>	Mar 09, 2023
<u>AB</u>	<u>!</u> NOVITIUM PHARMA	<u>200MG/5ML</u>	<u>A212252</u>	<u>001</u>	Jul 10, 2020
<u>AB</u>	RUBICON	<u>200MG/5ML</u>	<u>A215724</u>	<u>001</u>	Aug 18, 2022

TABLET; BUCCAL

SITAVIG

<u>+</u> !	LNHC	50MG	N203791	001	Apr 12, 2013
------------	------	------	---------	-----	--------------

TABLET; ORAL

ACYCLOVIR

<u>AB</u>	APOTEX INC	<u>400MG</u>	<u>A077309</u>	<u>001</u>	Sep 29, 2005
<u>AB</u>		<u>800MG</u>	<u>A077309</u>	<u>002</u>	Sep 29, 2005
<u>AB</u>	CADILA PHARMS LTD	<u>400MG</u>	<u>A202168</u>	<u>001</u>	Nov 15, 2013
<u>AB</u>		<u>800MG</u>	<u>A202168</u>	<u>002</u>	Nov 15, 2013
<u>AB</u>	CARLSBAD	<u>400MG</u>	<u>A075382</u>	<u>001</u>	Apr 30, 1999
<u>AB</u>		<u>800MG</u>	<u>A075382</u>	<u>002</u>	Apr 30, 1999
<u>AB</u>	HERITAGE	<u>400MG</u>	<u>A074891</u>	<u>001</u>	Oct 31, 1997
<u>AB</u>		<u>800MG</u>	<u>A074891</u>	<u>002</u>	Oct 31, 1997
<u>AB</u>	HETERO LABS LTD V	<u>400MG</u>	<u>A203834</u>	<u>001</u>	Oct 29, 2013
<u>AB</u>	<u>!</u>	<u>800MG</u>	<u>A203834</u>	<u>002</u>	Oct 29, 2013
<u>AB</u>	SQUARE PHARMS	<u>400MG</u>	<u>A209366</u>	<u>001</u>	Oct 07, 2019
<u>AB</u>		<u>800MG</u>	<u>A209366</u>	<u>002</u>	Oct 07, 2019
<u>AB</u>	STRIDES PHARMA	<u>400MG</u>	<u>A074946</u>	<u>001</u>	Nov 19, 1997
<u>AB</u>		<u>800MG</u>	<u>A074946</u>	<u>002</u>	Nov 19, 1997
<u>AB</u>	TEVA	<u>400MG</u>	<u>A074556</u>	<u>002</u>	Apr 22, 1997
<u>AB</u>		<u>800MG</u>	<u>A074556</u>	<u>003</u>	Apr 22, 1997
<u>AB</u>	YILING	<u>400MG</u>	<u>A210401</u>	<u>001</u>	Mar 07, 2018
<u>AB</u>		<u>800MG</u>	<u>A210401</u>	<u>002</u>	Mar 07, 2018
<u>AB</u>	ZYDUS PHARMS	<u>400MG</u>	<u>A204314</u>	<u>001</u>	Aug 19, 2014
<u>AB</u>		<u>800MG</u>	<u>A204314</u>	<u>002</u>	Aug 19, 2014

ACYCLOVIR SODIUM

INJECTABLE; INJECTION

ACYCLOVIR SODIUM

<u>AP</u>	EUGIA PHARMA	<u>EQ 50MG BASE/ML</u>	<u>A203701</u>	<u>001</u>	Oct 11, 2013
<u>AP</u>	<u>!</u> FRESENIUS KABI USA	<u>EQ 50MG BASE/ML</u>	<u>A074930</u>	<u>001</u>	May 13, 1998
<u>AP</u>		<u>EQ 50MG BASE/ML</u>	<u>A215404</u>	<u>001</u>	Jun 25, 2024
<u>AP</u>	SLATE RUN PHARMA	<u>EQ 50MG BASE/ML</u>	<u>A218111</u>	<u>001</u>	Jan 08, 2024
<u>AP</u>	ZYDUS PHARMS	<u>EQ 50MG BASE/ML</u>	<u>A206535</u>	<u>001</u>	Aug 31, 2018

ACYCLOVIR; HYDROCORTISONE

CREAM; TOPICAL

XERESE

<u>+</u> !	BAUSCH	5%; 1%	N022436	001	Jul 31, 2009
------------	--------	--------	---------	-----	--------------

ADAGRASIB

TABLET; ORAL

KRAZATI

<u>+</u> !	BRISTOL	200MG	N216340	001	Dec 12, 2022
------------	---------	-------	---------	-----	--------------

PRESCRIPTION DRUG PRODUCT LIST

ADAPALENE

CREAM; TOPICAL

ADAPALENE

AB	FOUGERA PHARMS	0.1%	A090824 001	Jun 30, 2010
-----------	----------------	-------------	--------------------	--------------

DIFFERIN

AB	+! GALDERMA LABS LP	0.1%	N020748 001	May 26, 2000
-----------	---------------------	-------------	--------------------	--------------

GEL; TOPICAL

ADAPALENE

AB	ACTAVIS MID ATLANTIC	0.3%	A201000 001	Oct 27, 2014
-----------	----------------------	-------------	--------------------	--------------

AB	ALEMBIC	0.3%	A213508 001	Jun 18, 2020
-----------	---------	-------------	--------------------	--------------

AB	ENCUBE ETHICALS	0.3%	A200298 001	Jun 14, 2012
-----------	-----------------	-------------	--------------------	--------------

AB	TARO	0.3%	A208322 001	Jun 23, 2016
-----------	------	-------------	--------------------	--------------

DIFFERIN

AB	+! GALDERMA LABS LP	0.3%	N021753 001	Jun 19, 2007
-----------	---------------------	-------------	--------------------	--------------

SOLUTION; TOPICAL

ADAPALENE

AB	CALL INC	0.1%	A203981 001	Sep 23, 2016
-----------	----------	-------------	--------------------	--------------

AB		0.1%	A204593 001	Jan 05, 2016
-----------	--	-------------	--------------------	--------------

ADAPALENE; BENZOYL PEROXIDE

GEL; TOPICAL

ADAPALENE AND BENZOYL PEROXIDE

AB	ACTAVIS LABS UT INC	0.3%;2.5%	A209641 001	Jun 22, 2022
-----------	---------------------	------------------	--------------------	--------------

AB	ALEMBIC	0.3%;2.5%	A214185 001	Aug 04, 2022
-----------	---------	------------------	--------------------	--------------

AB	ENCUBE	0.1%;2.5%	A206164 001	May 23, 2018
-----------	--------	------------------	--------------------	--------------

AB	GLENMARK PHARMS LTD	0.1%;2.5%	A208108 001	Nov 08, 2019
-----------	---------------------	------------------	--------------------	--------------

AB	PADAGIS ISRAEL	0.1%;2.5%	A205033 001	Jan 23, 2018
-----------	----------------	------------------	--------------------	--------------

AB		0.3%;2.5%	A212464 001	May 31, 2022
-----------	--	------------------	--------------------	--------------

AB	TARO	0.1%;2.5%	A206959 001	Jan 24, 2018
-----------	------	------------------	--------------------	--------------

AB		0.3%;2.5%	A209148 001	Oct 17, 2018
-----------	--	------------------	--------------------	--------------

AB	ZYDUS PHARMS	0.3%;2.5%	A214553 001	Jun 03, 2022
-----------	--------------	------------------	--------------------	--------------

EPIDUO

AB	+! GALDERMA LABS LP	0.1%;2.5%	N022320 001	Dec 08, 2008
-----------	---------------------	------------------	--------------------	--------------

EPIDUO FORTE

AB	+! GALDERMA LABS	0.3%;2.5%	N207917 001	Jul 15, 2015
-----------	------------------	------------------	--------------------	--------------

ADAPALENE; BENZOYL PEROXIDE; CLINDAMYCIN PHOSPHATE

GEL; TOPICAL

CABTREO

+!	BAUSCH	0.15%;3.1%;1.2%	N216632 001	Oct 20, 2023
----	--------	-----------------	-------------	--------------

ADEFOVIR DIPIVOXIL

TABLET; ORAL

ADEFOVIR DIPIVOXIL

AB	APOTEX	10MG	A205459 001	Jul 06, 2018
-----------	--------	-------------	--------------------	--------------

AB	! SIGMAPHARM LABS LLC	10MG	A202051 001	Aug 29, 2013
-----------	-----------------------	-------------	--------------------	--------------

ADENOSINE

INJECTABLE; INJECTION

ADENOSINE

AP	FRESENIUS KABI USA	3MG/ML	A077133 001	Apr 27, 2005
-----------	--------------------	---------------	--------------------	--------------

AP		3MG/ML	A205568 001	Apr 16, 2018
-----------	--	---------------	--------------------	--------------

AP	GLAND PHARMA LTD	3MG/ML	A077283 001	Jun 14, 2007
-----------	------------------	---------------	--------------------	--------------

AP		3MG/ML	A206778 001	Feb 16, 2018
-----------	--	---------------	--------------------	--------------

AP	HIKMA	3MG/ML	A076404 001	Jun 16, 2004
-----------	-------	---------------	--------------------	--------------

AP		3MG/ML	A076500 001	Jun 16, 2004
-----------	--	---------------	--------------------	--------------

AP	MYLAN LABS LTD	3MG/ML	A078686 001	May 13, 2009
-----------	----------------	---------------	--------------------	--------------

AP	! RISING	3MG/ML	A078076 001	Oct 31, 2008
-----------	----------	---------------	--------------------	--------------

SOLUTION; INTRAVENOUS

ADENOSINE

AP	AVET LIFESCIENCES	60MG/20ML (3MG/ML)	A202313 001	Sep 15, 2014
-----------	-------------------	---------------------------	--------------------	--------------

AP		90MG/30ML (3MG/ML)	A202313 002	Sep 15, 2014
-----------	--	---------------------------	--------------------	--------------

AP	EUGIA PHARMA	60MG/20ML (3MG/ML)	A205331 001	Nov 02, 2017
-----------	--------------	---------------------------	--------------------	--------------

AP		90MG/30ML (3MG/ML)	A205331 002	Nov 02, 2017
-----------	--	---------------------------	--------------------	--------------

AP	FRESENIUS KABI USA	60MG/20ML (3MG/ML)	A077897 001	Nov 27, 2017
-----------	--------------------	---------------------------	--------------------	--------------

AP		90MG/30ML (3MG/ML)	A077897 002	Nov 27, 2017
-----------	--	---------------------------	--------------------	--------------

AP	! MEITHEAL	60MG/20ML (3MG/ML)	A077425 001	Aug 29, 2013
-----------	------------	---------------------------	--------------------	--------------

AP	!	90MG/30ML (3MG/ML)	A077425 002	Aug 29, 2013
-----------	---	---------------------------	--------------------	--------------

AP	MYLAN ASI	60MG/20ML (3MG/ML)	A090212 001	Mar 28, 2014
-----------	-----------	---------------------------	--------------------	--------------

AP		90MG/30ML (3MG/ML)	A090212 002	Mar 28, 2014
-----------	--	---------------------------	--------------------	--------------

AP	RISING	60MG/20ML (3MG/ML)	A090450 001	Oct 02, 2014
-----------	--------	---------------------------	--------------------	--------------

AP		90MG/30ML (3MG/ML)	A090450 002	Oct 02, 2014
-----------	--	---------------------------	--------------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

AFAMELANOTIDE

IMPLANT; SUBCUTANEOUS

SCENESSE

+! CLIVUNEL INC

16MG

N210797 001 Oct 08, 2019

AFATINIB DIMALEATE

TABLET; ORAL

GILOTRIF

+ BOEHRINGER

EQ 20MG BASE

N201292 001 Jul 12, 2013

INGELHEIM

+

EQ 30MG BASE

N201292 002 Jul 12, 2013

+!

EQ 40MG BASE

N201292 003 Jul 12, 2013

AIR POLYMER-TYPE A

FOAM; INTRAUTERINE

EXEM FOAM KIT

+! GISKIT

10ML

N212279 001 Nov 07, 2019

ALBENDAZOLE

TABLET; ORAL

ALBENDAZOLEAB ACTAVIS ELIZABETH200MGA208094 001 May 20, 2019AB ALEMBIC200MGA215652 001 Sep 04, 2024AB ! DR REDDYS200MGA211034 001 Jan 26, 2021AB EDENBRIDGE PHARMS200MGA211117 001 May 14, 2019AB MSN200MGA213435 001 Jan 21, 2021AB ZYDUS PHARMS200MGA208979 001 Dec 14, 2018ALBUMIN HUMAN

INJECTABLE; INJECTION

OPTISON

+! GE HEALTHCARE

10MG/ML

N020899 001 Dec 31, 1997

ALBUTEROL SULFATE

AEROSOL, METERED; INHALATION

ALBUTEROL SULFATEAB1 CIPLAEQ 0.09MG BASE/INHA209959 001 Apr 08, 2020AB1 SANDOZEQ 0.09MG BASE/INHA207085 001 Jun 01, 2021PROVENTIL-HFAAB1 +! KINDEVAEQ 0.09MG BASE/INHN020503 001 Aug 15, 1996ALBUTEROL SULFATEAB2 ARMSTRONG PHARMSEQ 0.09MG BASE/INHA212447 001 May 21, 2024AB2 LUPINEQ 0.09MG BASE/INHA209954 001 Aug 24, 2020PROAIR HFAAB2 +! TEVA BRANDED PHARMEQ 0.09MG BASE/INHN021457 001 Oct 29, 2004

VENTOLIN HFA

BX +! GLAXOSMITHKLINE

EQ 0.09MG BASE/INH

N020983 001 Apr 19, 2001

POWDER, METERED; INHALATION

PROAIR RESPICLICK

+! TEVA BRANDED PHARM

EQ 0.09MG BASE/INH

N205636 001 Mar 31, 2015

SOLUTION; INHALATION

ALBUTEROL SULFATEAN LEXENPHARMEQ 0.021% BASEA215571 001 May 03, 2024ANEQ 0.042% BASEA215571 002 May 03, 2024AN LUOXIN AUROVITASEQ 0.021% BASEA211888 001 Apr 20, 2020ANEQ 0.042% BASEA211888 002 Apr 20, 2020ANEQ 0.083% BASEA206224 001 Oct 17, 2017AN NEPHRONEQ 0.021% BASEA076355 002 Mar 31, 2010AN !EQ 0.042% BASEA076355 001 Jun 28, 2004AN !EQ 0.083% BASEA074880 001 Sep 17, 1997AN !EQ 0.5% BASEA075664 001 Jun 26, 2001AN ! RITEDOSE CORPEQ 0.021% BASEA214531 001 Dec 28, 2021ANEQ 0.042% BASEA214531 002 Dec 28, 2021ANEQ 0.083% BASEA077839 001 Dec 16, 2008AN SENTISSEQ 0.5% BASEA074543 001 Jan 15, 1998AN SUN PHARMEQ 0.083% BASEA207857 001 Jul 21, 2017

SYRUP; ORAL

ALBUTEROL SULFATEAA AMNEAL PHARMSEQ 2MG BASE/5MLA079241 001 May 12, 2010AA CHARTWELL MOLECULAREQ 2MG BASE/5MLA078105 001 Dec 27, 2006AA CHARTWELL RXEQ 2MG BASE/5MLA077788 001 Jun 26, 2007AA COSETTEEQ 2MG BASE/5MLA074454 001 Sep 25, 1995AA QUAGENEQ 2MG BASE/5MLA212197 001 Sep 06, 2019AA ! TEVAEQ 2MG BASE/5MLA073419 001 Mar 30, 1992

PRESCRIPTION DRUG PRODUCT LIST

ALBUTEROL SULFATE

TABLET; ORAL

ALBUTEROL SULFATE

<u>AB</u>		AMNEAL PHARMS CO	<u>EQ 2MG BASE</u>	<u>A208804 001</u>	May 21, 2018
<u>AB</u>			<u>EQ 4MG BASE</u>	<u>A208804 002</u>	May 21, 2018
<u>AB</u>		DASH PHARMS NATCO	<u>EQ 2MG BASE</u>	<u>A072894 002</u>	Jan 17, 1991
<u>AB</u>	!		<u>EQ 4MG BASE</u>	<u>A072894 001</u>	Jan 17, 1991
<u>AB</u>		RISING	<u>EQ 2MG BASE</u>	<u>A207046 001</u>	Jun 29, 2018
<u>AB</u>			<u>EQ 4MG BASE</u>	<u>A207046 002</u>	Jun 29, 2018
<u>AB</u>		SUN PHARM INDUSTRIES	<u>EQ 2MG BASE</u>	<u>A072637 002</u>	Dec 05, 1989
<u>AB</u>			<u>EQ 4MG BASE</u>	<u>A072637 001</u>	Dec 05, 1989
<u>AB</u>		VIRTUS PHARM	<u>EQ 2MG BASE</u>	<u>A211397 001</u>	Oct 26, 2018
<u>AB</u>			<u>EQ 4MG BASE</u>	<u>A211397 002</u>	Oct 26, 2018
<u>AB</u>		ZYDUS PHARMS	<u>EQ 2MG BASE</u>	<u>A208884 001</u>	Oct 22, 2020
<u>AB</u>			<u>EQ 4MG BASE</u>	<u>A208884 002</u>	Oct 22, 2020
BX		AIZANT	EQ 2MG BASE	A210948 001	Mar 15, 2019
BX			EQ 4MG BASE	A210948 002	Mar 15, 2019

ALBUTEROL SULFATE; BUDESONIDE

AEROSOL, METERED; INHALATION

AIRSUPRA

+	!	ASTRAZENECA	EQ 0.09MG BASE/INH;0.08MG/INH	N214070 001	Jan 10, 2023
---	---	-------------	-------------------------------	-------------	--------------

ALBUTEROL SULFATE; IPRATROPIUM BROMIDE

SOLUTION; INHALATION

ALBUTEROL SULFATE AND IPRATROPIUM BROMIDE

<u>AN</u>		CIPLA	<u>EQ 0.083% BASE;0.017%</u>	<u>A077559 001</u>	Dec 31, 2007
<u>AN</u>	!	NEPHRON	<u>EQ 0.083% BASE;0.017%</u>	<u>A076749 001</u>	Dec 31, 2007
<u>AN</u>		RITEDOSE CORP	<u>EQ 0.083% BASE;0.017%</u>	<u>A202496 001</u>	Oct 01, 2012
<u>AN</u>		SUN PHARM	<u>EQ 0.083% BASE;0.017%</u>	<u>A207875 001</u>	Aug 07, 2017
		SPRAY, METERED; INHALATION COMBIVENT RESPIMAT			
	+	!	BOEHRINGER INGELHEIM	EQ 0.1MG BASE/INH;0.02MG/INH	N021747 001 Oct 07, 2011

ALCLOMETASONE DIPROPIONATE

CREAM; TOPICAL

ALCLOMETASONE DIPROPIONATE

<u>AB</u>	!	FOUGERA PHARMS	<u>0.05%</u>	<u>A076973 001</u>	Jul 12, 2005
<u>AB</u>		GLENMARK PHARMS LTD	<u>0.05%</u>	<u>A079061 001</u>	Jun 23, 2009
<u>AB</u>		TARO	<u>0.05%</u>	<u>A076587 001</u>	Sep 15, 2005

OINTMENT; TOPICAL

ALCLOMETASONE DIPROPIONATE

<u>AB</u>	!	FOUGERA PHARMS	<u>0.05%</u>	<u>A076884 001</u>	Jul 18, 2005
<u>AB</u>		GLENMARK PHARMS LTD	<u>0.05%</u>	<u>A079227 001</u>	Jul 30, 2009
<u>AB</u>		TARO	<u>0.05%</u>	<u>A076730 001</u>	Jul 29, 2004

ALCOHOL

SOLUTION; INTRA-ARTERIAL

ABLYSINOL

+	!	BPI LABS	99% (5ML)	N207987 002	Jun 21, 2018
---	---	----------	-----------	-------------	--------------

ALECTINIB HYDROCHLORIDE

CAPSULE; ORAL

ALECENSA

+	!	HOFFMANN-LA ROCHE	EQ 150MG BASE	N208434 001	Dec 11, 2015
---	---	-------------------	---------------	-------------	--------------

ALENDRONATE SODIUM

SOLUTION; ORAL

ALENDRONATE SODIUM

<u>AA</u>	!	HIKMA	<u>EQ 70MG BASE/75ML</u>	<u>A090520 001</u>	Feb 25, 2013
<u>AA</u>		NOVITIUM PHARMA	<u>EQ 70MG BASE/75ML</u>	<u>A214512 001</u>	May 11, 2023

TABLET; ORAL

ALENDRONATE SODIUM

<u>AB</u>		APOTEX	<u>EQ 5MG BASE</u>	<u>A077982 001</u>	Aug 04, 2008
<u>AB</u>			<u>EQ 10MG BASE</u>	<u>A077982 002</u>	Aug 04, 2008
<u>AB</u>			<u>EQ 35MG BASE</u>	<u>A077982 003</u>	Aug 04, 2008
<u>AB</u>			<u>EQ 70MG BASE</u>	<u>A077982 004</u>	Aug 04, 2008
<u>AB</u>		AUROBINDO PHARMA	<u>EQ 10MG BASE</u>	<u>A090124 001</u>	Aug 04, 2008
<u>AB</u>			<u>EQ 35MG BASE</u>	<u>A090124 002</u>	Aug 04, 2008
<u>AB</u>			<u>EQ 70MG BASE</u>	<u>A090124 003</u>	Aug 04, 2008
<u>AB</u>		CIPLA	<u>EQ 5MG BASE</u>	<u>A076768 001</u>	Aug 04, 2008
<u>AB</u>			<u>EQ 10MG BASE</u>	<u>A076768 002</u>	Aug 04, 2008
<u>AB</u>			<u>EQ 35MG BASE</u>	<u>A076768 003</u>	Aug 04, 2008
<u>AB</u>			<u>EQ 40MG BASE</u>	<u>A076768 004</u>	Aug 04, 2008

PRESCRIPTION DRUG PRODUCT LIST

ALENDRONATE SODIUM

TABLET; ORAL

ALENDRONATE SODIUM

AB		<u>EQ 70MG BASE</u>	<u>A076768 005</u>	Aug 04, 2008
AB	HANGZHOU BINJIANG	<u>EQ 5MG BASE</u>	<u>A090258 001</u>	Sep 24, 2009
AB		<u>EQ 10MG BASE</u>	<u>A090258 002</u>	Sep 24, 2009
AB		<u>EQ 35MG BASE</u>	<u>A090258 003</u>	Sep 24, 2009
AB		<u>EQ 70MG BASE</u>	<u>A090258 004</u>	Sep 24, 2009
AB	IPCA LABS LTD	<u>EQ 10MG BASE</u>	<u>A206387 001</u>	Jun 04, 2024
AB		<u>EQ 35MG BASE</u>	<u>A206387 002</u>	Jun 04, 2024
AB		<u>EQ 40MG BASE</u>	<u>A206387 003</u>	Jun 04, 2024
AB		<u>EQ 70MG BASE</u>	<u>A206387 004</u>	Jun 04, 2024
AB	SUN PHARM	<u>EQ 5MG BASE</u>	<u>A090022 001</u>	Sep 10, 2008
AB		<u>EQ 10MG BASE</u>	<u>A090022 002</u>	Sep 10, 2008
AB		<u>EQ 35MG BASE</u>	<u>A090022 003</u>	Sep 10, 2008
AB		<u>EQ 70MG BASE</u>	<u>A090022 004</u>	Sep 10, 2008
AB	WATSON LABS	<u>EQ 35MG BASE</u>	<u>A076984 001</u>	Aug 04, 2008
AB		<u>EQ 40MG BASE</u>	<u>A076984 002</u>	Aug 04, 2008
AB		<u>EQ 70MG BASE</u>	<u>A076984 003</u>	Aug 04, 2008

FOSAMAX

AB	+! ORGANON	<u>EQ 70MG BASE</u>	<u>N020560 005</u>	Oct 20, 2000
	TABLET, EFFERVESCENT; ORAL			
	BINOSTO			
	+! RADIUS	<u>EQ 70MG BASE</u>	<u>N202344 001</u>	Mar 12, 2012

ALENDRONATE SODIUM; CHOLECALCIFEROL

TABLET; ORAL

FOSAMAX PLUS D

	+ ORGANON LLC	<u>EQ 70MG BASE; 2,800 IU</u>	<u>N021762 001</u>	Apr 07, 2005
	+!	<u>EQ 70MG BASE; 5,600 IU</u>	<u>N021762 002</u>	Apr 26, 2007

ALFENTANIL HYDROCHLORIDE

INJECTABLE; INJECTION

ALFENTA

AP	+! RISING	<u>EQ 0.5MG BASE/ML</u>	<u>N019353 001</u>	Dec 29, 1986
	<u>ALFENTANIL</u>			
AP	HOSPIRA	<u>EQ 0.5MG BASE/ML</u>	<u>A075221 001</u>	Oct 28, 1999

ALFUZOSIN HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL

ALFUZOSIN HYDROCHLORIDE

AB	APOTEX INC	<u>10MG</u>	<u>A079013 001</u>	Jul 18, 2011
AB	AUROBINDO PHARMA LTD	<u>10MG</u>	<u>A079060 001</u>	Aug 30, 2012
AB	INVAGEN PHARMS	<u>10MG</u>	<u>A090284 001</u>	Jan 17, 2012
AB	UNICHEM	<u>10MG</u>	<u>A203192 001</u>	Jan 28, 2016
	<u>UROXATRAL</u>			
AB	+! ADVANZ PHARMA	<u>10MG</u>	<u>N021287 001</u>	Jun 12, 2003

ALISKIREN HEMIFUMARATE

TABLET; ORAL

ALISKIREN HEMIFUMARATE

AB	ENDO OPERATIONS	<u>EQ 150MG BASE</u>	<u>A206665 001</u>	Mar 22, 2019
AB		<u>EQ 300MG BASE</u>	<u>A206665 002</u>	Mar 22, 2019
	<u>TEKTURNA</u>			
AB	+ NODEN PHARMA	<u>EQ 150MG BASE</u>	<u>N021985 001</u>	Mar 05, 2007
AB	+!	<u>EQ 300MG BASE</u>	<u>N021985 002</u>	Mar 05, 2007

ALITRETINOLIN

GEL; TOPICAL

PANRETIN

	+! ADVANZ PHARMA	<u>EQ 0.1% BASE</u>	<u>N020886 001</u>	Feb 02, 1999
--	------------------	----------------------------	---------------------------	--------------

ALLOPURINOL

TABLET; ORAL

ALLOPURINOL

AB	ACCORD HLTHCARE	<u>100MG</u>	<u>A203154 001</u>	May 06, 2013
AB		<u>300MG</u>	<u>A203154 002</u>	May 06, 2013
AB	CHARTWELL	<u>100MG</u>	<u>A077353 001</u>	Sep 08, 2005
AB		<u>300MG</u>	<u>A077353 002</u>	Sep 08, 2005
AB	ENDO OPERATIONS	<u>100MG</u>	<u>A075798 001</u>	Jun 27, 2003
AB		<u>300MG</u>	<u>A075798 002</u>	Jun 27, 2003
AB	HARMAN FINOCHEM	<u>100MG</u>	<u>A214443 001</u>	Mar 07, 2022
AB		<u>200MG</u>	<u>A214443 003</u>	Jun 17, 2024
AB		<u>300MG</u>	<u>A214443 002</u>	Mar 07, 2022
AB	HETERO LABS LTD V	<u>100MG</u>	<u>A217748 001</u>	Aug 03, 2023

PRESCRIPTION DRUG PRODUCT LIST

ALLOPURINOL

TABLET; ORAL

ALLOPURINOL

<u>AB</u>		<u>300MG</u>	<u>A217748 002</u>	Aug 03, 2023
<u>AB</u>	INDOCO	<u>100MG</u>	<u>A204467 001</u>	Jul 28, 2016
<u>AB</u>		<u>300MG</u>	<u>A204467 002</u>	Jul 28, 2016
<u>AB</u>	MYLAN	<u>100MG</u>	<u>A018659 001</u>	Oct 24, 1986
<u>AB</u>		<u>300MG</u>	<u>A018659 002</u>	Oct 24, 1986
<u>AB</u>	NORTHSTAR HLTHCARE	<u>100MG</u>	<u>A078253 001</u>	Sep 11, 2007
<u>AB</u>		<u>300MG</u>	<u>A078253 002</u>	Sep 11, 2007
<u>AB</u>	SUN PHARM INDUSTRIES	<u>100MG</u>	<u>A071450 002</u>	Jan 09, 1987
<u>AB</u>		<u>300MG</u>	<u>A071450 001</u>	Jan 09, 1987
<u>AB</u>	UNICHEM	<u>100MG</u>	<u>A211820 001</u>	Mar 12, 2019
<u>AB</u>	!	<u>300MG</u>	<u>A211820 002</u>	Mar 12, 2019
<u>AB</u>	WATSON LABS	<u>100MG</u>	<u>N018832 002</u>	Sep 28, 1984
<u>AB</u>		<u>300MG</u>	<u>N018877 001</u>	Sep 28, 1984
<u>AB</u>	ZYDUS PHARMS	<u>100MG</u>	<u>A210117 001</u>	Oct 12, 2017
<u>AB</u>		<u>300MG</u>	<u>A210117 002</u>	Oct 12, 2017
<u>LOPURIN</u>				
<u>AB</u>	DR REDDYS	<u>100MG</u>	<u>A071586 001</u>	Apr 02, 1987
<u>AB</u>		<u>300MG</u>	<u>A071587 001</u>	Apr 02, 1987
<u>ZYLOPRIM</u>				
<u>AB</u>	+ CASPER PHARMA LLC	<u>100MG</u>	<u>N016084 001</u>	
<u>AB</u>	+	<u>200MG</u>	<u>N016084 003</u>	Aug 04, 2022
<u>AB</u>	+	<u>300MG</u>	<u>N016084 002</u>	

ALLOPURINOL SODIUM

INJECTABLE; INJECTION

ALLOPURINOL SODIUM

<u>AP</u>	GLAND PHARMA LTD	<u>EQ 500MG BASE/VIAL</u>	<u>A212363 001</u>	Jan 26, 2022
<u>AP</u>	HIKMA	<u>EQ 500MG BASE/VIAL</u>	<u>A076870 001</u>	Aug 26, 2004
<u>ALOPRIM</u>				
<u>AP</u>	+! MYLAN	<u>EQ 500MG BASE/VIAL</u>	<u>N020298 001</u>	May 17, 1996

ALMOTRIPTAN MALATE

TABLET; ORAL

ALMOTRIPTAN MALATE

<u>AB</u>	AJANTA PHARMA LTD	<u>EQ 6.25MG BASE</u>	<u>A205523 001</u>	Mar 03, 2016
<u>AB</u>	!	<u>EQ 12.5MG BASE</u>	<u>A205523 002</u>	Mar 03, 2016
<u>AB</u>	MYLAN	<u>EQ 6.25MG BASE</u>	<u>A205171 001</u>	Nov 09, 2015
<u>AB</u>		<u>EQ 12.5MG BASE</u>	<u>A205171 002</u>	Nov 09, 2015
<u>AB</u>	TEVA PHARMS USA	<u>EQ 6.25MG BASE</u>	<u>A078027 001</u>	Jul 07, 2015
<u>AB</u>		<u>EQ 12.5MG BASE</u>	<u>A078027 002</u>	Jul 07, 2015

ALOGLIPTIN BENZOATE

TABLET; ORAL

NESINA

+	TAKEDA PHARMS USA	EQ 6.25MG BASE	N022271 001	Jan 25, 2013
+		EQ 12.5MG BASE	N022271 002	Jan 25, 2013
+	!	EQ 25MG BASE	N022271 003	Jan 25, 2013

ALOGLIPTIN BENZOATE; METFORMIN HYDROCHLORIDE

TABLET; ORAL

KAZANO

+	TAKEDA PHARMS USA	EQ 12.5MG BASE;500MG	N203414 001	Jan 25, 2013
+	!	EQ 12.5MG BASE;1GM	N203414 002	Jan 25, 2013

ALOGLIPTIN BENZOATE; PIOGLITAZONE HYDROCHLORIDE

TABLET; ORAL

OSEN

+	TAKEDA PHARMS USA	EQ 12.5MG BASE;EQ 30MG BASE	N022426 005	Jan 25, 2013
+		EQ 25MG BASE;EQ 15MG BASE	N022426 001	Jan 25, 2013
+		EQ 25MG BASE;EQ 30MG BASE	N022426 002	Jan 25, 2013
+	!	EQ 25MG BASE;EQ 45MG BASE	N022426 003	Jan 25, 2013

ALOSETRON HYDROCHLORIDE

TABLET; ORAL

ALOSETRON HYDROCHLORIDE

<u>AB</u>	AMNEAL PHARMS	<u>EQ 0.5MG BASE</u>	<u>A206647 001</u>	Dec 22, 2016
<u>AB</u>		<u>EQ 1MG BASE</u>	<u>A206647 002</u>	Dec 22, 2016
<u>AB</u>	MANKIND PHARMA	<u>EQ 0.5MG BASE</u>	<u>A213614 001</u>	Sep 09, 2020
<u>AB</u>		<u>EQ 1MG BASE</u>	<u>A213614 002</u>	Sep 09, 2020
<u>AB</u>	RISING	<u>EQ 0.5MG BASE</u>	<u>A209180 001</u>	Jan 14, 2019
<u>AB</u>		<u>EQ 1MG BASE</u>	<u>A209180 002</u>	Jan 14, 2019

PRESCRIPTION DRUG PRODUCT LIST

ALOSETRON HYDROCHLORIDE

TABLET; ORAL

LOTRONEX

AB	+	LEGACY PHARMA	EQ 0.5MG BASE	N021107 002	Dec 23, 2003
AB	+	!	EQ 1MG BASE	N021107 001	Feb 09, 2000

ALPELISIB

GRANULES; ORAL

VIJOICE

+	NOVARTIS	50MG/PACKET	N218466 001	Apr 24, 2024
---	----------	-------------	-------------	--------------

TABLET; ORAL

PIQRAY

+	NOVARTIS	50MG	N212526 001	May 24, 2019
---	----------	------	-------------	--------------

+		150MG	N212526 002	May 24, 2019
---	--	-------	-------------	--------------

+	!	200MG	N212526 003	May 24, 2019
---	---	-------	-------------	--------------

VIJOICE

+	NOVARTIS	50MG	N215039 001	Apr 05, 2022
---	----------	------	-------------	--------------

+		125MG	N215039 002	Apr 05, 2022
---	--	-------	-------------	--------------

+	!	200MG	N215039 003	Apr 05, 2022
---	---	-------	-------------	--------------

ALPHA-TOCOPHEROL ACETATE; ASCORBIC ACID; BIOTIN; CHOLECALCIFEROL; CYANOCOBALAMIN; DEXPANTHENOL; FOLIC ACID; NIACINAMIDE; PYRIDOXINE HYDROCHLORIDE; RIBOFLAVIN 5'-PHOSPHATE SODIUM; THIAMINE HYDROCHLORIDE; VITAMIN A PALMITATE; VITAMIN K

SOLUTION; INTRAVENOUS

INFUVITE ADULT

+	SANDOZ CANADA INC	2 IU/ML; 40MG/ML; 12MCG/ML; 40 IU/ML; 1MCG/ML; 3MG/ML; 120MCG/ML; 8MG/ML; 1.2MG/ML; 0.72MG/ML; 1.2MG/ML; 660 IU/ML; 0.03MG/ML	N021163 001	May 18, 2000
---	-------------------	---	-------------	--------------

+	!	2 IU/ML; 40MG/ML; 12MCG/ML; 40 IU/ML; 1MCG/ML; 3MG/ML; 120MCG/ML; 8MG/ML; 1.2MG/ML; 0.72MG/ML; 1.2MG/ML; 660 IU/ML; 30MCG/ML	N021163 002	Jun 16, 2003
---	---	--	-------------	--------------

ALPRAZOLAM

CONCENTRATE; ORAL

ALPRAZOLAM

!	HIKMA	1MG/ML	A074312 001	Oct 31, 1993
---	-------	--------	-------------	--------------

TABLET; ORAL

ALPRAZOLAM

AB		ACTAVIS ELIZABETH	0.25MG	A074342 001	Oct 31, 1993
AB			0.5MG	A074342 002	Oct 31, 1993
AB			1MG	A074342 003	Oct 31, 1993
AB			2MG	A074342 004	Oct 31, 1993
AB		APOTEX INC	0.25MG	A077741 001	Jan 19, 2007
AB			0.5MG	A077741 002	Jan 19, 2007
AB			1MG	A077741 003	Jan 19, 2007
AB			2MG	A077741 004	Jan 19, 2007
AB		AUROBINDO PHARMA	0.25MG	A203346 001	Jul 31, 2015
AB			0.5MG	A203346 002	Jul 31, 2015
AB			1MG	A203346 003	Jul 31, 2015
AB			2MG	A203346 004	Jul 31, 2015
AB		CHARTWELL RX	0.25MG	A207507 001	Jul 09, 2018
AB			0.5MG	A207507 002	Jul 09, 2018
AB			1MG	A207507 003	Jul 09, 2018
AB			2MG	A207507 004	Jul 09, 2018
AB		NATCO	0.25MG	A200739 001	Apr 15, 2015
AB			0.5MG	A200739 002	Apr 15, 2015
AB			1MG	A200739 003	Apr 15, 2015
AB			2MG	A200739 004	Apr 15, 2015
AB		NOVITIUM PHARMA	0.25MG	A074174 001	Oct 19, 1993
AB			0.5MG	A074174 002	Oct 19, 1993
AB			1MG	A074174 003	Oct 19, 1993
AB			2MG	A074174 004	Oct 19, 1993
AB		SANDOZ	0.25MG	A074112 001	Dec 29, 1995
AB			0.5MG	A074112 002	Dec 29, 1995
AB			1MG	A074112 003	Dec 29, 1995
AB			2MG	A074909 001	Mar 25, 1998
AB		STRIDES PHARMA	0.25MG	A090248 001	Sep 17, 2010
AB			0.5MG	A090248 002	Sep 17, 2010
AB			1MG	A090248 003	Sep 17, 2010
AB			2MG	A090248 004	Sep 17, 2010
AB		SUN PHARM	0.25MG	A090082 001	Jun 17, 2010
AB			0.5MG	A090082 002	Jun 17, 2010
AB			1MG	A090082 003	Jun 17, 2010

PRESCRIPTION DRUG PRODUCT LIST

ALPRAZOLAM

TABLET; ORAL

ALPRAZOLAM

<u>AB</u>		<u>2MG</u>	<u>A090082 004</u>	Jun 17, 2010
	<u>XANAX</u>			
<u>AB</u>	+	UPJOHN	<u>0.25MG</u>	<u>N018276 001</u>
<u>AB</u>	+		<u>0.5MG</u>	<u>N018276 002</u>
<u>AB</u>	+	!	<u>1MG</u>	<u>N018276 003</u>
<u>AB</u>	+		<u>2MG</u>	<u>N018276 004</u>

TABLET, EXTENDED RELEASE; ORAL

ALPRAZOLAM

<u>AB</u>		ACTAVIS ELIZABETH	<u>0.5MG</u>	<u>A078056 001</u>	Feb 13, 2007
<u>AB</u>			<u>1MG</u>	<u>A078056 002</u>	Feb 13, 2007
<u>AB</u>			<u>2MG</u>	<u>A078056 003</u>	Feb 13, 2007
<u>AB</u>			<u>3MG</u>	<u>A078056 004</u>	Feb 13, 2007
<u>AB</u>		AMNEAL PHARMS NY	<u>0.5MG</u>	<u>A078387 001</u>	May 30, 2008
<u>AB</u>			<u>1MG</u>	<u>A078387 002</u>	May 30, 2008
<u>AB</u>			<u>2MG</u>	<u>A078387 003</u>	May 30, 2008
<u>AB</u>			<u>3MG</u>	<u>A078387 004</u>	May 30, 2008
<u>AB</u>		APOTEX INC	<u>0.5MG</u>	<u>A078449 001</u>	Nov 12, 2008
<u>AB</u>			<u>1MG</u>	<u>A078449 004</u>	Dec 23, 2015
<u>AB</u>			<u>2MG</u>	<u>A078449 002</u>	Nov 12, 2008
<u>AB</u>			<u>3MG</u>	<u>A078449 003</u>	Nov 12, 2008
<u>AB</u>		AUROBINDO PHARMA	<u>0.5MG</u>	<u>A090871 001</u>	Jun 07, 2011
<u>AB</u>			<u>1MG</u>	<u>A090871 002</u>	Jun 07, 2011
<u>AB</u>			<u>2MG</u>	<u>A090871 003</u>	Jun 07, 2011
<u>AB</u>			<u>3MG</u>	<u>A090871 004</u>	Jun 07, 2011
	<u>XANAX XR</u>				
<u>AB</u>	+	UPJOHN	<u>0.5MG</u>	<u>N021434 001</u>	Jan 17, 2003
<u>AB</u>	+		<u>1MG</u>	<u>N021434 002</u>	Jan 17, 2003
<u>AB</u>	+		<u>2MG</u>	<u>N021434 003</u>	Jan 17, 2003
<u>AB</u>	+	!	<u>3MG</u>	<u>N021434 004</u>	Jan 17, 2003

TABLET, ORALLY DISINTEGRATING; ORAL

ALPRAZOLAM

<u>AB</u>		ACTAVIS ELIZABETH	<u>0.25MG</u>	<u>A078561 001</u>	Mar 16, 2010
<u>AB</u>			<u>0.5MG</u>	<u>A078561 002</u>	Mar 16, 2010
<u>AB</u>			<u>1MG</u>	<u>A078561 003</u>	Mar 16, 2010
<u>AB</u>			<u>2MG</u>	<u>A078561 004</u>	Mar 16, 2010
<u>AB</u>		ENDO OPERATIONS	<u>0.25MG</u>	<u>A078088 001</u>	Jan 09, 2009
<u>AB</u>			<u>0.5MG</u>	<u>A078088 002</u>	Jan 09, 2009
<u>AB</u>	!		<u>1MG</u>	<u>A078088 003</u>	Jan 09, 2009
<u>AB</u>			<u>2MG</u>	<u>A078088 004</u>	Jan 09, 2009

ALPROSTADIL

INJECTABLE; INJECTION

ALPROSTADIL

<u>AP</u>		HIKMA	<u>0.5MG/ML</u>	<u>A074815 001</u>	Jan 20, 1998
<u>AP</u>		MEITHEAL	<u>0.5MG/ML</u>	<u>A075196 001</u>	Apr 30, 1999
	<u>CAVERJECT</u>				
<u>AP</u>	+	PFIZER	<u>0.01MG/VIAL</u>	<u>N020379 001</u>	Jul 06, 1995
<u>AP</u>	+	!	<u>0.02MG/VIAL</u>	<u>N020379 002</u>	Jul 06, 1995
<u>AP</u>	+	!	<u>0.04MG/VIAL</u>	<u>N020379 004</u>	May 19, 1997
	<u>EDEX</u>				
<u>AP</u>	+	ENDO OPERATIONS	<u>0.01MG/VIAL</u>	<u>N020649 002</u>	Jun 12, 1997
<u>AP</u>	+		<u>0.02MG/VIAL</u>	<u>N020649 003</u>	Jun 12, 1997
<u>AP</u>	+	!	<u>0.04MG/VIAL</u>	<u>N020649 004</u>	Jun 12, 1997
	<u>PROSTIN VR PEDIATRIC</u>				
<u>AP</u>	+	PFIZER	<u>0.5MG/ML</u>	<u>N018484 001</u>	
	CAVERJECT IMPULSE				
		PFIZER	0.01MG/VIAL	N021212 001	Jun 11, 2002
			0.02MG/VIAL	N021212 002	Jun 11, 2002
	EDEX				
	+	ENDO OPERATIONS	0.01MG/VIAL	N020649 005	Jul 30, 1998
	+		0.02MG/VIAL	N020649 006	Jul 30, 1998
	+		0.04MG/VIAL	N020649 007	Jul 30, 1998

ALVIMOPAN

CAPSULE; ORAL

ALVIMOPAN

<u>AB</u>		ENDO OPERATIONS	<u>12MG</u>	<u>A216843 001</u>	Jan 24, 2023
<u>AB</u>		HIKMA	<u>12MG</u>	<u>A217753 001</u>	Aug 31, 2023
<u>AB</u>	!	WATSON LABS TEVA	<u>12MG</u>	<u>A208295 001</u>	Dec 19, 2019

PRESCRIPTION DRUG PRODUCT LIST

AMANTADINE HYDROCHLORIDE

CAPSULE; ORAL

AMANTADINE HYDROCHLORIDE

<u>AB</u>		ALEMBIC	<u>100MG</u>	<u>A208966</u>	<u>001</u>	Jun 21, 2017
<u>AB</u>	!	BIONPHARMA	<u>100MG</u>	<u>A078720</u>	<u>001</u>	May 29, 2008
<u>AB</u>		HERITAGE PHARMA	<u>100MG</u>	<u>A209171</u>	<u>001</u>	Jun 12, 2017
<u>AB</u>		HUMANWELL PURACAP	<u>100MG</u>	<u>A214580</u>	<u>001</u>	Dec 20, 2022
<u>AB</u>		RISING	<u>100MG</u>	<u>A210129</u>	<u>001</u>	Mar 02, 2020
<u>AB</u>		RUBICON	<u>100MG</u>	<u>A212044</u>	<u>001</u>	May 21, 2020
<u>AB</u>		SANDOZ	<u>100MG</u>	<u>A071293</u>	<u>001</u>	Feb 18, 1987
<u>AB</u>		STRIDES PHARMA	<u>100MG</u>	<u>A209047</u>	<u>001</u>	Jun 07, 2017
<u>AB</u>		STRIDES SOFTGELS	<u>100MG</u>	<u>A211354</u>	<u>001</u>	Feb 18, 2022
<u>AB</u>		UPSHER SMITH LABS	<u>100MG</u>	<u>A070589</u>	<u>001</u>	Aug 05, 1986
<u>AB</u>		ZYDUS PHARMS	<u>100MG</u>	<u>A208278</u>	<u>001</u>	May 31, 2016

CAPSULE, EXTENDED RELEASE; ORAL

AMANTADINE HYDROCHLORIDE

<u>AB</u>		ZYDUS	<u>EQ 68.5MG BASE</u>	<u>A214897</u>	<u>001</u>	Aug 26, 2024
<u>GOCOVRI</u>						
<u>AB</u>	+	SUPERNUS PHARMS	<u>EQ 68.5MG BASE</u>	<u>N208944</u>	<u>001</u>	Aug 24, 2017
	+	!	EQ 137MG BASE	N208944	002	Aug 24, 2017

SYRUP; ORAL

AMANTADINE HYDROCHLORIDE

<u>AA</u>	!	AUROBINDO PHARMA USA	<u>50MG/5ML</u>	<u>A074170</u>	<u>001</u>	Oct 28, 1994
<u>AA</u>	!	CHARTWELL RX	<u>50MG/5ML</u>	<u>A074028</u>	<u>001</u>	Jun 28, 1993
<u>AA</u>		ELYSIUM	<u>50MG/5ML</u>	<u>A214178</u>	<u>001</u>	Aug 20, 2021
<u>AA</u>	!	PHARM ASSOC	<u>50MG/5ML</u>	<u>A074509</u>	<u>001</u>	Jul 17, 1995
<u>AA</u>		XTRTRIUM LABS INC	<u>50MG/5ML</u>	<u>A075060</u>	<u>001</u>	Dec 24, 1998

TABLET; ORAL

AMANTADINE HYDROCHLORIDE

<u>AB</u>		ALEMBIC	<u>100MG</u>	<u>A214284</u>	<u>001</u>	Oct 15, 2020
<u>AB</u>		ATHEM	<u>100MG</u>	<u>A210215</u>	<u>001</u>	Mar 10, 2020
<u>AB</u>		STRIDES PHARMA	<u>100MG</u>	<u>A209035</u>	<u>001</u>	Jun 09, 2017
<u>AB</u>	!	UPSHER SMITH LABS	<u>100MG</u>	<u>A076186</u>	<u>001</u>	Dec 16, 2002
<u>AB</u>		WATSON LABS INC	<u>100MG</u>	<u>A208096</u>	<u>001</u>	Dec 15, 2016
<u>AB</u>		ZHEJIANG JUTAI PHARM	<u>100MG</u>	<u>A218598</u>	<u>001</u>	Sep 26, 2024

AMBRISENTAN

TABLET; ORAL

AMBRISENTAN

<u>AB</u>		APOTEX	<u>5MG</u>	<u>A210701</u>	<u>001</u>	May 19, 2022
<u>AB</u>			<u>10MG</u>	<u>A210701</u>	<u>002</u>	May 19, 2022
<u>AB</u>		AUROBINDO PHARMA	<u>5MG</u>	<u>A216531</u>	<u>001</u>	Jul 21, 2022
<u>AB</u>			<u>10MG</u>	<u>A216531</u>	<u>002</u>	Jul 21, 2022
<u>AB</u>		CIPLA	<u>5MG</u>	<u>A210715</u>	<u>001</u>	Apr 26, 2019
<u>AB</u>			<u>10MG</u>	<u>A210715</u>	<u>002</u>	Apr 26, 2019
<u>AB</u>		MYLAN	<u>5MG</u>	<u>A208441</u>	<u>001</u>	Mar 28, 2019
<u>AB</u>			<u>10MG</u>	<u>A208441</u>	<u>002</u>	Mar 28, 2019
<u>AB</u>		SIGMAPHARM LABS LLC	<u>5MG</u>	<u>A208354</u>	<u>001</u>	Apr 10, 2019
<u>AB</u>			<u>10MG</u>	<u>A208354</u>	<u>002</u>	Apr 10, 2019
<u>AB</u>		SUN PHARM	<u>5MG</u>	<u>A210784</u>	<u>001</u>	Mar 28, 2019
<u>AB</u>			<u>10MG</u>	<u>A210784</u>	<u>002</u>	Mar 28, 2019
<u>AB</u>		WATSON LABS INC	<u>5MG</u>	<u>A208252</u>	<u>001</u>	Mar 28, 2019
<u>AB</u>			<u>10MG</u>	<u>A208252</u>	<u>002</u>	Mar 28, 2019
<u>AB</u>		ZYDUS PHARMS	<u>5MG</u>	<u>A210058</u>	<u>001</u>	Mar 28, 2019
<u>AB</u>			<u>10MG</u>	<u>A210058</u>	<u>002</u>	Mar 28, 2019

LETAIRIS

<u>AB</u>	+	GILEAD	<u>5MG</u>	<u>N022081</u>	<u>001</u>	Jun 15, 2007
<u>AB</u>	+	!	<u>10MG</u>	<u>N022081</u>	<u>002</u>	Jun 15, 2007

AMCINONIDE

CREAM; TOPICAL

AMCINONIDE

<u>AB</u>	!	GENUS LIFESCIENCES	<u>0.1%</u>	<u>A076065</u>	<u>001</u>	May 15, 2003
<u>AB</u>		TARO PHARM INDS	<u>0.1%</u>	<u>A076229</u>	<u>001</u>	May 31, 2002

LOTION; TOPICAL

AMCINONIDE

!

OINTMENT; TOPICAL

		GENUS	0.1%	A076329	001	Nov 06, 2002
<u>AMCINONIDE</u>						
<u>AB</u>	!	GENUS	<u>0.1%</u>	<u>A076096</u>	<u>001</u>	Nov 19, 2002
<u>AB</u>		TARO PHARM INDS	<u>0.1%</u>	<u>A076367</u>	<u>001</u>	Mar 19, 2003

PRESCRIPTION DRUG PRODUCT LIST

AMIFAMPRIDINE PHOSPHATE

TABLET; ORAL

FIRDAPSE

+! CATALYST PHARMS EQ 10MG BASE N208078 001 Nov 28, 2018

AMIKACIN SULFATE

INJECTABLE; INJECTION

AMIKACIN SULFATE**AP** ! AVET LIFESCIENCES **EQ 250MG BASE/ML** **A204040 001** Dec 12, 2013**AP** FRESENIUS KABI USA **EQ 250MG BASE/ML** **A205604 001** Dec 09, 2015**AP** HIKMA **EQ 250MG BASE/ML** **A063315 001** Apr 11, 1994**AP** MEITHEAL **EQ 250MG BASE/ML** **A064045 002** Sep 28, 1993**AP** QILU **EQ 250MG BASE/ML** **A218146 001** Mar 19, 2024**AP** SAGENT PHARMS INC **EQ 250MG BASE/ML** **A203323 001** May 12, 2016

! HIKMA EQ 50MG BASE/ML A063313 001 Apr 11, 1994

SUSPENSION, LIPOSOMAL; INHALATION

ARIKAYCE KIT

+! INSMED INC EQ 590MG BASE/8.4ML N207356 001 Sep 28, 2018

AMILORIDE HYDROCHLORIDE

TABLET; ORAL

AMILORIDE HYDROCHLORIDE**AB** ! ENDO OPERATIONS **5MG** **A070346 001** Jan 22, 1986**AB** SIGMAPHARM LABS LLC **5MG** **A079133 001** Jan 30, 2009MIDAMOR**AB** + PADAGIS US **5MG** **N018200 001**AMILORIDE HYDROCHLORIDE; HYDROCHLOROTHIAZIDE

TABLET; ORAL

AMILORIDE HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE**AB** BARR **EQ 5MG ANHYDROUS; 50MG** **A071111 001** May 10, 1988**AB** ! RISING **EQ 5MG ANHYDROUS; 50MG** **A073209 001** Oct 31, 1991AMINO ACIDS

INJECTABLE; INJECTION

AMINO ACIDS

B BRAUN 15% (150GM/1000ML) A091112 001 Apr 13, 2012

15% (300GM/2000ML) A091112 002 Apr 13, 2012

AMINOSYN II 10% IN PLASTIC CONTAINER

ICU MEDICAL INC 10% (10GM/100ML) N020015 001 Dec 19, 1991

AMINOSYN II 15% IN PLASTIC CONTAINER

ICU MEDICAL INC 15% (15GM/100ML) N020041 001 Dec 19, 1991

AMINOSYN-PF 10%

ICU MEDICAL INC 10% (10GM/100ML) N019492 002 Oct 17, 1986

AMINOSYN-PF 7%

ICU MEDICAL INC 7% (7GM/100ML) N019398 001 Sep 06, 1985

CLINISOL 15% SULFITE FREE IN PLASTIC CONTAINER

BAXTER HLTHCARE 15% (15GM/100ML) A020512 001 Aug 30, 1996

PREMASOL 10% IN PLASTIC CONTAINER

BAXTER HLTHCARE 10% (10GM/100ML) A075880 002 Jun 19, 2003

PREMASOL 6% IN PLASTIC CONTAINER

BAXTER HLTHCARE 6% (6GM/100ML) A075880 001 Jun 19, 2003

PROSOL 20% SULFITE FREE IN PLASTIC CONTAINER

+! BAXTER HLTHCARE 20% (20GM/100ML) N020849 001 Aug 26, 1998

TRAVASOL 10% IN PLASTIC CONTAINER

BAXTER HLTHCARE 10% (10MG/100ML) N018931 003 Aug 23, 1984

TRAVASOL 5.5% IN PLASTIC CONTAINER

BAXTER HLTHCARE 5.5% (5.5GM/100ML) N018931 001 Aug 23, 1984

TRAVASOL 8.5% IN PLASTIC CONTAINER

BAXTER HLTHCARE 8.5% (8.5GM/100ML) N018931 002 Aug 23, 1984

TROPHAMINE

+! B BRAUN 6% (6GM/100ML) N019018 001 Jul 20, 1984

TROPHAMINE 10%

+! B BRAUN 10% (10GM/100ML) N019018 003 Sep 07, 1988

AMINO ACIDS; CALCIUM CHLORIDE; DEXTROSE; MAGNESIUM CHLORIDE; POTASSIUM PHOSPHATE, DIBASIC; SODIUM ACETATE; SODIUM CHLORIDE

INJECTABLE; INJECTION

CLINIMIX E 2.75/10 SULFITE FREE W/ ELECT IN DEXTROSE 10% W/ CALCIUM IN PLASTIC CONTAINER

+! BAXTER HLTHCARE 2.75%; 33MG/100ML; 10GM/100ML; 51MG/100ML; 261MG/100ML; 217MG/100ML; 112MG/100ML N020678 002 Mar 26, 1997

CLINIMIX E 2.75/25 SULFITE FREE W/ ELECT IN DEXTROSE 25% W/ CALCIUM IN PLASTIC CONTAINER

+! BAXTER HLTHCARE 2.75%; 33MG/100ML; 25GM/100ML; 51MG/100ML; 261MG/100ML; 217MG/100ML; 112MG/100ML N020678 005 Mar 26, 1997

PRESCRIPTION DRUG PRODUCT LIST

AMINO ACIDS; CALCIUM CHLORIDE; DEXTROSE; MAGNESIUM CHLORIDE; POTASSIUM PHOSPHATE, DIBASIC; SODIUM ACETATE; SODIUM CHLORIDE

INJECTABLE; INJECTION

CLINIMIX E 2.75/5 SULFITE FREE W/ ELECT IN DEXTROSE 5% W/ CALCIUM IN PLASTIC CONTAINER				
+! BAXTER HLTHCARE	2.75%;33MG/100ML;5GM/100ML;51MG/100ML;2	N020678 001	Mar 26,	1997
	61MG/100ML;217MG/100ML;112MG/100ML			
CLINIMIX E 4.25/10 SULFITE FREE W/ ELECT IN DEXTROSE 10% W/ CALCIUM IN PLASTIC CONTAINER				
+! BAXTER HLTHCARE	4.25%;33MG/100ML;10GM/100ML;51MG/100ML;	N020678 009	Mar 26,	1997
	261MG/100ML;297MG/100ML;77MG/100ML			
CLINIMIX E 4.25/20 SULFITE FREE W/ ELECT IN DEXTROSE 20% W/ CALCIUM IN PLASTIC CONTAINER				
+! BAXTER HLTHCARE	4.25%;33MG/100ML;20GM/100ML;51MG/100ML;	N020678 011	Mar 26,	1997
	261MG/100ML;297MG/100ML;77MG/100ML			
CLINIMIX E 4.25/25 SULFITE FREE W/ ELECT IN DEXTROSE 25% W/ CALCIUM IN PLASTIC CONTAINER				
+! BAXTER HLTHCARE	4.25%;33MG/100ML;25GM/100ML;51MG/100ML;	N020678 012	Mar 26,	1997
	261MG/100ML;297MG/100ML;77MG/100ML			
CLINIMIX E 4.25/5 SULFITE FREE W/ ELECT IN DEXTROSE 5% W/ CALCIUM IN PLASTIC CONTAINER				
+! BAXTER HLTHCARE	4.25%;33MG/100ML;5GM/100ML;51MG/100ML;2	N020678 008	Mar 26,	1997
	61MG/100ML;297MG/100ML;77MG/100ML			
CLINIMIX E 5/10 SULFITE FREE W/ ELECT IN DEXTROSE 10% W/ CALCIUM IN PLASTIC CONTAINER				
+! BAXTER HLTHCARE	5%;33MG/100ML;10GM/100ML;51MG/100ML;261	N020678 016	Mar 26,	1997
	MG/100ML;340MG/100ML;59MG/100ML			
CLINIMIX E 5/15 SULFITE FREE W/ ELECT IN DEXTROSE 15% W/ CALCIUM IN PLASTIC CONTAINER				
+! BAXTER HLTHCARE	5%;33MG/100ML;15GM/100ML;51MG/100ML;261	N020678 017	Mar 26,	1997
	MG/100ML;340MG/100ML;59MG/100ML			
CLINIMIX E 5/20 SULFITE FREE W/ ELECT IN 20% DEXTROSE W/ CALCIUM IN PLASTIC CONTAINER				
+! BAXTER HLTHCARE	5%;33MG/100ML;20GM/100ML;51MG/100ML;261	N020678 018	Mar 26,	1997
	MG/100ML;340MG/100ML;59MG/100ML			
CLINIMIX E 5/25 SULFITE FREE W/ ELECT IN DEXTROSE 25% W/ CALCIUM IN PLASTIC CONTAINER				
+! BAXTER HLTHCARE	5%;33MG/100ML;25GM/100ML;51MG/100ML;261	N020678 019	Mar 26,	1997
	MG/100ML;340MG/100ML;59MG/100ML			
CLINIMIX E 5/35 SULFITE FREE W/ ELECT IN DEXTROSE 35% W/ CALCIUM IN PLASTIC CONTAINER				
+! BAXTER HLTHCARE	5%;33MG/100ML;35GM/100ML;51MG/100ML;261	N020678 021	Mar 26,	1997
	MG/100ML;340MG/100ML;59MG/100ML			

AMINO ACIDS; CALCIUM CHLORIDE; DEXTROSE; MAGNESIUM SULFATE; POTASSIUM CHLORIDE; SODIUM ACETATE; SODIUM GLYCEROPHOSPHATE; SOYBEAN OIL

EMULSION; INTRAVENOUS

KABIVEN IN PLASTIC CONTAINER				
+ FRESENIUS KABI USA	3.3%;29MG/100ML;9.8GM/100ML;96MG/100ML;	N200656 004	Aug 25,	2014
	174MG/100ML;239MG/100ML			
	;147MG/100ML;3.9GM/100ML (1026ML)			
+ FRESENIUS KABI USA	3.3%;29MG/100ML;9.8GM/100ML;96MG/100ML;	N200656 005	Aug 25,	2014
	174MG/100ML;239MG/100ML;147MG/100ML;3.9			
	GM/100ML (1540ML)			
+ FRESENIUS KABI USA	3.3%;29MG/100ML;9.8GM/100ML;96MG/100ML;	N200656 006	Aug 25,	2014
	174MG/100ML;239MG/100ML;147MG/100ML;3.9			
	GM/100ML (2053ML)			
+! FRESENIUS KABI USA	3.3%;29MG/100ML;9.8GM/100ML;96MG/100ML;	N200656 007	Aug 25,	2014
	174MG/100ML;239MG/100ML;147MG/100ML;3.9			
	GM/100ML (2566ML)			
PERIKABIVEN IN PLASTIC CONTAINER				
+ FRESENIUS KABI USA	2.4%;20MG/100ML;6.8GM/100ML;68MG/100ML;	N200656 001	Aug 25,	2014
	124MG/100ML;170MG/100ML;105MG/100ML;3.5			
	GM/100ML (1440ML)			
+ FRESENIUS KABI USA	2.4%;20MG/100ML;6.8GM/100ML;68MG/100ML;	N200656 002	Aug 25,	2014
	124MG/100ML;170MG/100ML;105MG/100ML;3.5			
	GM/100ML (1920ML)			
+! FRESENIUS KABI USA	2.4%;20MG/100ML;6.8GM/100ML;68MG/100ML;	N200656 003	Aug 25,	2014
	124MG/100ML;170MG/100ML			
	;105MG/100ML;3.5GM/100ML (2400ML)			

AMINO ACIDS; DEXTROSE

INJECTABLE; INJECTION

CLINIMIX 2.75/10 SULFITE FREE IN DEXTROSE 10% IN PLASTIC CONTAINER				
BAXTER HLTHCARE	2.75%;10GM/100ML	N020734 002	Sep 29,	1997
CLINIMIX 2.75/25 SULFITE FREE IN DEXTROSE 25% IN PLASTIC CONTAINER				
BAXTER HLTHCARE	2.75%;25GM/100ML	N020734 005	Sep 29,	1997
CLINIMIX 2.75/5 SULFITE FREE IN DEXTROSE 5% IN PLASTIC CONTAINER				
BAXTER HLTHCARE	2.75%;5GM/100ML	N020734 001	Sep 29,	1997
CLINIMIX 4.25/10 SULFITE FREE IN DEXTROSE 10% IN PLASTIC CONTAINER				
BAXTER HLTHCARE	4.25%;10GM/100ML	N020734 008	Sep 29,	1997
CLINIMIX 4.25/20 SULFITE FREE IN DEXTROSE 20% IN PLASTIC CONTAINER				
BAXTER HLTHCARE	4.25%;20GM/100ML	N020734 010	Sep 29,	1997
CLINIMIX 4.25/25 SULFITE FREE IN DEXTROSE 25% IN PLASTIC CONTAINER				
BAXTER HLTHCARE	4.25%;25GM/100ML	N020734 011	Sep 29,	1997
CLINIMIX 4.25/5 SULFITE FREE IN DEXTROSE 5% IN PLASTIC CONTAINER				
BAXTER HLTHCARE	4.25%;5GM/100ML	N020734 007	Sep 29,	1997

PRESCRIPTION DRUG PRODUCT LIST

AMINO ACIDS; DEXTROSE

INJECTABLE; INJECTION

CLINIMIX 5/10 SULFITE FREE	IN DEXTROSE 10%	IN PLASTIC CONTAINER			
BAXTER HLTHCARE	5%;10GM/100ML		N020734	014	Sep 29, 1997
CLINIMIX 5/15 SULFITE FREE	IN DEXTROSE 15%	IN PLASTIC CONTAINER			
BAXTER HLTHCARE	5%;15GM/100ML		N020734	015	Sep 29, 1997
CLINIMIX 5/20 SULFITE FREE	IN DEXTROSE 20%	IN PLASTIC CONTAINER			
BAXTER HLTHCARE	5%;20GM/100ML		N020734	016	Sep 29, 1997
CLINIMIX 5/25 SULFITE FREE	IN DEXTROSE 25%	IN PLASTIC CONTAINER			
BAXTER HLTHCARE	5%;25GM/100ML		N020734	017	Sep 29, 1997
CLINIMIX 5/35 SULFITE FREE	IN DEXTROSE 35%	IN PLASTIC CONTAINER			
BAXTER HLTHCARE	5%;35GM/100ML		N020734	018	Sep 29, 1997
CLINIMIX 8/14 SULFITE FREE	IN DEXTROSE 14%	IN PLASTIC CONTAINER			
BAXTER HLTHCARE	8%;14GM/100ML		N020734	019	Apr 13, 2021

AMINOCAPROIC ACID

INJECTABLE; INJECTION

AMINOCAPROIC ACID

AP LUITPOLD **250MG/ML** **A071192 001** Dec 01, 1987

AMINOCAPROIC ACID IN PLASTIC CONTAINER

AP ! HOSPIRA **250MG/ML** **A070010 001** Mar 09, 1987

SOLUTION; ORAL

AMICAR

AA + HIKMA **0.25GM/ML** **N015230 002**

AMINOCAPROIC ACID

AA AMNEAL **0.25GM/ML** **A212780 001** Aug 23, 2019
AA ANNORA PHARMA **0.25GM/ML** **A216464 001** Nov 04, 2022
AA AUROBINDO PHARMA **0.25GM/ML** **A216804 001** Sep 26, 2022
AA BELCHER **0.25GM/ML** **A213825 001** Apr 08, 2021
AA CARNEGIE **0.25GM/ML** **A214140 001** Jan 26, 2021
AA FLORIDA **0.25GM/ML** **A215510 001** Mar 23, 2022
AA MSN **0.25GM/ML** **A215100 001** Sep 26, 2024
AA SUNNY **0.25GM/ML** **A213213 001** Dec 27, 2023
AA TARO **0.25GM/ML** **A214458 001** Mar 24, 2023
AA ! TRUPHARMA **0.25GM/ML** **A212494 001** Aug 11, 2020

TABLET; ORAL

AMICAR

AB + HIKMA **500MG** **N015197 001**

AB + **1GM** **N015197 002** Jun 24, 2004

AMINOCAPROIC ACID

AB AMNEAL **500MG** **A212492 001** Nov 26, 2019
AB APPCO **500MG** **A213944 001** Sep 14, 2022
AB **1GM** **A213944 002** Sep 14, 2022
AB CARNEGIE **500MG** **A213928 001** Feb 12, 2021
AB **1GM** **A213928 002** Feb 12, 2021
AB MSN **500MG** **A212938 001** Nov 06, 2020
AB **1GM** **A212938 002** Sep 13, 2023
AB SUNNY **500MG** **A209060 001** Nov 27, 2018
AB ! **1GM** **A209060 002** Nov 27, 2018

AMINOLEVULINIC ACID HYDROCHLORIDE

FOR SOLUTION; ORAL

GLEOLAN

+! NXDC 1.5GM/VIAL N208630 001 Jun 06, 2017

GEL; TOPICAL

AMELUZ

+! BIOFRONTERA 10% N208081 001 May 10, 2016

SOLUTION; TOPICAL

LEVULAN

+! SUN PHARM INDS INC 20% N020965 001 Dec 03, 1999

AMINOPHYLLINE

INJECTABLE; INJECTION

AMINOPHYLLINE

! HOSPIRA 25MG/ML A087242 001 Oct 26, 1983

AMINOSALICYLIC ACID

GRANULE, DELAYED RELEASE; ORAL

PASER

! EUROCEPT PHARMS 4GM/PACKET A074346 001 Jun 30, 1994

PRESCRIPTION DRUG PRODUCT LIST

AMIODARONE HYDROCHLORIDE

INJECTABLE; INJECTION

AMIODARONE HYDROCHLORIDE

<u>AP</u>	ACELLA	<u>50MG/ML</u>	<u>A077610 001</u>	Oct 30, 2008	
<u>AP</u>		<u>50MG/ML</u>	<u>A077834 001</u>	Oct 30, 2008	
<u>AP</u>	!	FRESENIUS KABI USA	<u>50MG/ML</u>	<u>A075761 001</u>	Oct 15, 2002
<u>AP</u>	!	GLAND PHARMA LTD	<u>50MG/ML</u>	<u>A077161 001</u>	Apr 20, 2005
<u>AP</u>		HIKMA FARMACEUTICA	<u>50MG/ML</u>	<u>A077234 001</u>	Feb 25, 2008
<u>AP</u>	!	MYLAN INSTITUTIONAL	<u>50MG/ML</u>	<u>A076217 001</u>	Oct 15, 2002
<u>AP</u>		ZHEJIANG POLY PHARM	<u>50MG/ML</u>	<u>A218253 001</u>	Sep 11, 2024

NEXTERONE

+	!	BAXTER HLTHCARE	150MG/100ML (1.5MG/ML)	N022325 002	Nov 16, 2010
+	!		360MG/200ML (1.8MG/ML)	N022325 003	Nov 16, 2010

TABLET; ORAL

AMIODARONE HYDROCHLORIDE

<u>AB</u>		AUROBINDO PHARMA LTD	<u>200MG</u>	<u>A204742 001</u>	Jun 03, 2016
<u>AB</u>		CHARTWELL RX	<u>100MG</u>	<u>A077069 003</u>	Oct 04, 2016
<u>AB</u>			<u>200MG</u>	<u>A077069 001</u>	Apr 08, 2005
<u>AB</u>			<u>400MG</u>	<u>A077069 002</u>	Apr 08, 2005
<u>AB</u>		DR REDDYS LABS SA	<u>100MG</u>	<u>A075389 002</u>	Dec 28, 2017
<u>AB</u>			<u>200MG</u>	<u>A075389 001</u>	Jan 25, 2001
<u>AB</u>			<u>400MG</u>	<u>A075389 003</u>	Dec 28, 2017
<u>AB</u>		RUBICON	<u>100MG</u>	<u>A078578 002</u>	Feb 26, 2021
<u>AB</u>			<u>200MG</u>	<u>A078578 001</u>	Nov 06, 2008
<u>AB</u>			<u>400MG</u>	<u>A078578 003</u>	Feb 26, 2021
<u>AB</u>		TARO	<u>100MG</u>	<u>A075424 002</u>	Dec 18, 2002
<u>AB</u>	!		<u>200MG</u>	<u>A075424 001</u>	Mar 30, 2001
<u>AB</u>			<u>400MG</u>	<u>A076362 001</u>	Nov 29, 2002
<u>AB</u>		TEVA PHARMS	<u>200MG</u>	<u>A074739 001</u>	Nov 30, 1998
<u>AB</u>		UNICHEM	<u>200MG</u>	<u>A213446 001</u>	Jul 21, 2020
<u>AB</u>		ZYDUS PHARMS USA INC	<u>100MG</u>	<u>A079029 002</u>	Jan 06, 2023
<u>AB</u>			<u>200MG</u>	<u>A079029 001</u>	Sep 16, 2008
<u>AB</u>			<u>400MG</u>	<u>A079029 003</u>	Jan 06, 2023

PACERONE

<u>AB</u>		UPSHER SMITH LABS	<u>100MG</u>	<u>A075135 002</u>	Apr 12, 2005
<u>AB</u>			<u>200MG</u>	<u>A075135 001</u>	Apr 30, 1998
<u>AB</u>			<u>400MG</u>	<u>A075135 003</u>	Jul 02, 2020

AMIODARONE HYDROCHLORIDE

		TARO	300MG	A076362 002	Dec 02, 2003
--	--	------	-------	-------------	--------------

AMISULPRIDE

SOLUTION; INTRAVENOUS

BARHEMSYS

+	!	ACACIA	5MG/2ML (2.5MG/ML)	N209510 001	Feb 26, 2020
+	!		10MG/4ML (2.5MG/ML)	N209510 002	Sep 01, 2020

AMITRIPTYLINE HYDROCHLORIDE

TABLET; ORAL

AMITRIPTYLINE HYDROCHLORIDE

<u>AB</u>		ACCORD HLTHCARE	<u>10MG</u>	<u>A202446 001</u>	Jun 04, 2014
<u>AB</u>			<u>25MG</u>	<u>A202446 002</u>	Jun 04, 2014
<u>AB</u>			<u>50MG</u>	<u>A202446 003</u>	Jun 04, 2014
<u>AB</u>			<u>75MG</u>	<u>A202446 004</u>	Jun 04, 2014
<u>AB</u>			<u>100MG</u>	<u>A202446 005</u>	Jun 04, 2014
<u>AB</u>			<u>150MG</u>	<u>A202446 006</u>	Jun 04, 2014
<u>AB</u>		AIPING PHARM INC	<u>10MG</u>	<u>A212654 002</u>	Sep 29, 2021
<u>AB</u>			<u>25MG</u>	<u>A212654 001</u>	Apr 07, 2020
<u>AB</u>			<u>50MG</u>	<u>A212654 003</u>	Sep 29, 2021
<u>AB</u>			<u>75MG</u>	<u>A212654 004</u>	Sep 29, 2021
<u>AB</u>			<u>100MG</u>	<u>A212654 005</u>	Sep 29, 2021
<u>AB</u>			<u>150MG</u>	<u>A212654 006</u>	Sep 29, 2021
<u>AB</u>		BRECKENRIDGE	<u>10MG</u>	<u>A216243 001</u>	Jun 06, 2022
<u>AB</u>			<u>25MG</u>	<u>A216243 002</u>	Jun 06, 2022
<u>AB</u>			<u>50MG</u>	<u>A216243 003</u>	Jun 06, 2022
<u>AB</u>			<u>75MG</u>	<u>A216243 004</u>	Jun 06, 2022
<u>AB</u>			<u>100MG</u>	<u>A216243 005</u>	Jun 06, 2022
<u>AB</u>			<u>150MG</u>	<u>A216243 006</u>	Jun 06, 2022
<u>AB</u>		MANKIND PHARMA	<u>10MG</u>	<u>A213999 001</u>	Feb 19, 2021
<u>AB</u>			<u>25MG</u>	<u>A213999 002</u>	Feb 19, 2021
<u>AB</u>			<u>50MG</u>	<u>A213999 003</u>	Feb 19, 2021
<u>AB</u>			<u>75MG</u>	<u>A213999 004</u>	Feb 19, 2021
<u>AB</u>			<u>100MG</u>	<u>A213999 005</u>	Feb 19, 2021

PRESCRIPTION DRUG PRODUCT LIST

AMITRIPTYLINE HYDROCHLORIDE

TABLET; ORAL

AMITRIPTYLINE HYDROCHLORIDE

<u>AB</u>		<u>150MG</u>	<u>A213999 006</u>	Feb 19, 2021
<u>AB</u>	RISING	<u>10MG</u>	<u>A217411 001</u>	May 19, 2023
<u>AB</u>		<u>25MG</u>	<u>A217411 002</u>	May 19, 2023
<u>AB</u>		<u>50MG</u>	<u>A217411 003</u>	May 19, 2023
<u>AB</u>		<u>75MG</u>	<u>A217411 004</u>	May 19, 2023
<u>AB</u>		<u>100MG</u>	<u>A217411 005</u>	May 19, 2023
<u>AB</u>		<u>150MG</u>	<u>A217411 006</u>	May 19, 2023
<u>AB</u>	RUBICON	<u>10MG</u>	<u>A215376 001</u>	May 01, 2023
<u>AB</u>		<u>25MG</u>	<u>A215376 002</u>	May 01, 2023
<u>AB</u>		<u>50MG</u>	<u>A215376 003</u>	May 01, 2023
<u>AB</u>		<u>75MG</u>	<u>A215376 004</u>	May 01, 2023
<u>AB</u>		<u>100MG</u>	<u>A215376 005</u>	May 01, 2023
<u>AB</u>		<u>150MG</u>	<u>A215376 006</u>	May 01, 2023
<u>AB</u>	+ SANDOZ	<u>10MG</u>	<u>A085968 004</u>	
<u>AB</u>	+!	<u>25MG</u>	<u>A085968 002</u>	
<u>AB</u>	+	<u>50MG</u>	<u>A085968 001</u>	
<u>AB</u>	+	<u>75MG</u>	<u>A085968 006</u>	
<u>AB</u>	+	<u>100MG</u>	<u>A085968 003</u>	
<u>AB</u>	+	<u>150MG</u>	<u>A085968 005</u>	
<u>AB</u>	SUN PHARM INDS INC	<u>10MG</u>	<u>A089399 002</u>	Jul 14, 1987
<u>AB</u>		<u>25MG</u>	<u>A089399 001</u>	Jul 14, 1987
<u>AB</u>		<u>50MG</u>	<u>A089399 003</u>	Jul 14, 1987
<u>AB</u>		<u>75MG</u>	<u>A089399 004</u>	Jul 14, 1987
<u>AB</u>		<u>100MG</u>	<u>A089399 005</u>	Jul 14, 1987
<u>AB</u>		<u>150MG</u>	<u>A089399 006</u>	Jul 14, 1987
<u>AB</u>	UNICHEM	<u>10MG</u>	<u>A214548 001</u>	May 19, 2021
<u>AB</u>		<u>25MG</u>	<u>A214548 002</u>	May 19, 2021
<u>AB</u>		<u>50MG</u>	<u>A214548 003</u>	May 19, 2021
<u>AB</u>		<u>75MG</u>	<u>A214548 004</u>	May 19, 2021
<u>AB</u>		<u>100MG</u>	<u>A214548 005</u>	May 19, 2021
<u>AB</u>		<u>150MG</u>	<u>A214548 006</u>	May 19, 2021
<u>AB</u>	ZYDUS PHARMS	<u>10MG</u>	<u>A210086 001</u>	Oct 06, 2017
<u>AB</u>		<u>25MG</u>	<u>A210086 002</u>	Oct 06, 2017
<u>AB</u>		<u>50MG</u>	<u>A210086 003</u>	Oct 06, 2017
<u>AB</u>		<u>75MG</u>	<u>A210086 004</u>	Oct 06, 2017
<u>AB</u>		<u>100MG</u>	<u>A210086 005</u>	Oct 06, 2017
<u>AB</u>		<u>150MG</u>	<u>A210086 006</u>	Oct 06, 2017

AMITRIPTYLINE HYDROCHLORIDE; CHLORDIAZEPOXIDE

TABLET; ORAL

CHLORDIAZEPOXIDE AND AMITRIPTYLINE HYDROCHLORIDE

<u>AB</u>	MICRO LABS	<u>EQ 12.5MG BASE;5MG</u>	<u>A211925 001</u>	Feb 02, 2022
<u>AB</u>		<u>EQ 25MG BASE;10MG</u>	<u>A211925 002</u>	Feb 02, 2022
<u>AB</u>	MYLAN PHARMS INC	<u>EQ 12.5MG BASE;5MG</u>	<u>A071297 002</u>	Dec 10, 1986
<u>AB</u>	!	<u>EQ 25MG BASE;10MG</u>	<u>A071297 001</u>	Dec 10, 1986

AMITRIPTYLINE HYDROCHLORIDE; PERPHENAZINE

TABLET; ORAL

PERPHENAZINE AND AMITRIPTYLINE HYDROCHLORIDE

	MYLAN	10MG;2MG	A071443 002	Nov 10, 1988
		10MG;4MG	A071443 003	Nov 10, 1988
	!	25MG;2MG	A071443 004	Nov 10, 1988
	!	25MG;4MG	A071443 005	Nov 10, 1988
	!	50MG;4MG	A071443 001	Nov 10, 1988

AMLODIPINE BENZOATE

SUSPENSION; ORAL

KATERZIA

	+! AZURITY	EQ 1MG BASE/ML	N211340 001	Jul 08, 2019
--	------------	----------------	-------------	--------------

AMLODIPINE BESYLATE

SOLUTION; ORAL

NORLIQVA

	+! CMP DEV LLC	EQ 1MG BASE/ML	N214439 001	Feb 24, 2022
--	----------------	----------------	-------------	--------------

TABLET; ORAL

AMLODIPINE BESYLATE

<u>AB</u>	ACCORD HLTHCARE	<u>EQ 2.5MG BASE</u>	<u>A202553 001</u>	Apr 29, 2013
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A202553 002</u>	Apr 29, 2013
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A202553 003</u>	Apr 29, 2013
<u>AB</u>	ALKEM	<u>EQ 2.5MG BASE</u>	<u>A078925 001</u>	May 04, 2009
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A078925 002</u>	May 04, 2009

PRESCRIPTION DRUG PRODUCT LIST

AMLODIPINE BESYLATE

TABLET; ORAL

AMLODIPINE BESYLATE

<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A078925 003</u>	May 04, 2009
<u>AB</u>	AUROBINDO PHARMA	<u>EQ 2.5MG BASE</u>	<u>A078021 001</u>	Jul 17, 2007
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A078021 002</u>	Jul 17, 2007
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A078021 003</u>	Jul 17, 2007
<u>AB</u>	CHARTWELL RX	<u>EQ 2.5MG BASE</u>	<u>A076692 001</u>	Jul 20, 2007
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A076692 002</u>	Jul 20, 2007
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A076692 003</u>	Jul 20, 2007
<u>AB</u>	CHINA RESOURCES	<u>EQ 2.5MG BASE</u>	<u>A090752 003</u>	May 16, 2016
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A090752 001</u>	Apr 15, 2011
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A090752 002</u>	Apr 15, 2011
<u>AB</u>	CIPLA	<u>EQ 2.5MG BASE</u>	<u>A077073 001</u>	Sep 26, 2007
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A077073 002</u>	Sep 26, 2007
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A077073 003</u>	Sep 26, 2007
<u>AB</u>	COREPHARMA	<u>EQ 2.5MG BASE</u>	<u>A076719 001</u>	May 23, 2007
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A076719 002</u>	May 23, 2007
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A076719 003</u>	May 23, 2007
<u>AB</u>	EPIC PHARMA LLC	<u>EQ 2.5MG BASE</u>	<u>A078552 001</u>	Apr 08, 2009
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A078552 002</u>	Apr 08, 2009
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A078552 003</u>	Apr 08, 2009
<u>AB</u>	INVAGEN PHARMS	<u>EQ 2.5MG BASE</u>	<u>A077955 001</u>	Aug 28, 2007
<u>AB</u>		<u>EQ 2.5MG BASE</u>	<u>A206367 001</u>	Dec 10, 2015
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A077955 002</u>	Aug 28, 2007
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A206367 002</u>	Dec 10, 2015
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A077955 003</u>	Aug 28, 2007
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A206367 003</u>	Dec 10, 2015
<u>AB</u>	LUPIN	<u>EQ 2.5MG BASE</u>	<u>A078043 001</u>	Jul 12, 2007
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A078043 002</u>	Jul 12, 2007
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A078043 003</u>	Jul 12, 2007
<u>AB</u>	ORBION PHARMS	<u>EQ 2.5MG BASE</u>	<u>A078453 001</u>	Jul 02, 2009
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A078453 002</u>	Jul 02, 2009
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A078453 003</u>	Jul 02, 2009
<u>AB</u>	OXFORD PHARMS	<u>EQ 2.5MG BASE</u>	<u>A078414 001</u>	Apr 07, 2010
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A078414 002</u>	Apr 07, 2010
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A078414 003</u>	Apr 07, 2010
<u>AB</u>	POLYGEN PHARMS	<u>EQ 2.5MG BASE</u>	<u>A207821 001</u>	Jul 11, 2016
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A207821 002</u>	Jul 11, 2016
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A207821 003</u>	Jul 11, 2016
<u>AB</u>	STRIDES PHARMA	<u>EQ 2.5MG BASE</u>	<u>A077516 001</u>	Jul 11, 2007
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A077516 002</u>	Jul 11, 2007
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A077516 003</u>	Jul 11, 2007
<u>AB</u>	TEVA	<u>EQ 2.5MG BASE</u>	<u>A076846 001</u>	Jun 28, 2007
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A076846 002</u>	Jun 28, 2007
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A076846 003</u>	Jun 28, 2007
<u>AB</u>	UNICHEM	<u>EQ 2.5MG BASE</u>	<u>A203245 001</u>	Oct 21, 2013
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A203245 002</u>	Oct 21, 2013
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A203245 003</u>	Oct 21, 2013
<u>AB</u>	ZYDUS PHARMS USA	<u>EQ 2.5MG BASE</u>	<u>A078226 001</u>	Jul 09, 2007
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A078226 002</u>	Jul 09, 2007
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A078226 003</u>	Jul 09, 2007
<u>NORVASC</u>				
<u>AB</u>	+ VIATRIS	<u>EQ 2.5MG BASE</u>	<u>N019787 001</u>	Jul 31, 1992
<u>AB</u>	+	<u>EQ 5MG BASE</u>	<u>N019787 002</u>	Jul 31, 1992
<u>AB</u>	+!	<u>EQ 10MG BASE</u>	<u>N019787 003</u>	Jul 31, 1992

AMLODIPINE BESYLATE; ATORVASTATIN CALCIUM

TABLET; ORAL

AMLODIPINE BESYLATE AND ATORVASTATIN CALCIUM

<u>AB</u>	APOTEX	<u>EQ 5MG BASE;EQ 10MG BASE</u>	<u>A205199 001</u>	Nov 18, 2019
<u>AB</u>		<u>EQ 5MG BASE;EQ 20MG BASE</u>	<u>A205199 002</u>	Nov 18, 2019
<u>AB</u>		<u>EQ 5MG BASE;EQ 40MG BASE</u>	<u>A205199 003</u>	Nov 18, 2019
<u>AB</u>		<u>EQ 5MG BASE;EQ 80MG BASE</u>	<u>A205199 004</u>	Nov 18, 2019
<u>AB</u>		<u>EQ 10MG BASE;EQ 10MG BASE</u>	<u>A205199 005</u>	Nov 18, 2019
<u>AB</u>		<u>EQ 10MG BASE;EQ 20MG BASE</u>	<u>A205199 006</u>	Nov 18, 2019
<u>AB</u>		<u>EQ 10MG BASE;EQ 40MG BASE</u>	<u>A205199 007</u>	Nov 18, 2019
<u>AB</u>		<u>EQ 10MG BASE;EQ 80MG BASE</u>	<u>A205199 008</u>	Nov 18, 2019
<u>AB</u>	DR REDDYS	<u>EQ 2.5MG BASE;EQ 10MG BASE</u>	<u>A203874 001</u>	Mar 07, 2014
<u>AB</u>		<u>EQ 2.5MG BASE;EQ 20MG BASE</u>	<u>A203874 002</u>	Mar 07, 2014
<u>AB</u>		<u>EQ 2.5MG BASE;EQ 40MG BASE</u>	<u>A203874 003</u>	Mar 07, 2014
<u>AB</u>		<u>EQ 5MG BASE;EQ 10MG BASE</u>	<u>A203874 004</u>	Mar 07, 2014
<u>AB</u>		<u>EQ 5MG BASE;EQ 20MG BASE</u>	<u>A203874 005</u>	Mar 07, 2014

PRESCRIPTION DRUG PRODUCT LIST

AMLODIPINE BESYLATE; ATORVASTATIN CALCIUM

TABLET; ORAL

AMLODIPINE BESYLATE AND ATORVASTATIN CALCIUM

AB		<u>EQ 5MG BASE;EQ 40MG BASE</u>	<u>A203874 006</u>	Mar 07, 2014
AB		<u>EQ 5MG BASE;EQ 80MG BASE</u>	<u>A203874 007</u>	Mar 07, 2014
AB		<u>EQ 10MG BASE;EQ 10MG BASE</u>	<u>A203874 008</u>	Mar 07, 2014
AB		<u>EQ 10MG BASE;EQ 20MG BASE</u>	<u>A203874 009</u>	Mar 07, 2014
AB		<u>EQ 10MG BASE;EQ 40MG BASE</u>	<u>A203874 010</u>	Mar 07, 2014
AB		<u>EQ 10MG BASE;EQ 80MG BASE</u>	<u>A203874 011</u>	Mar 07, 2014
AB	MYLAN	<u>EQ 5MG BASE;EQ 10MG BASE</u>	<u>A200465 004</u>	Nov 29, 2013
AB		<u>EQ 5MG BASE;EQ 20MG BASE</u>	<u>A200465 005</u>	Nov 29, 2013
AB		<u>EQ 5MG BASE;EQ 40MG BASE</u>	<u>A200465 006</u>	Nov 29, 2013
AB		<u>EQ 5MG BASE;EQ 80MG BASE</u>	<u>A200465 007</u>	Nov 29, 2013
AB		<u>EQ 10MG BASE;EQ 10MG BASE</u>	<u>A200465 008</u>	Nov 29, 2013
AB		<u>EQ 10MG BASE;EQ 20MG BASE</u>	<u>A200465 009</u>	Nov 29, 2013
AB		<u>EQ 10MG BASE;EQ 40MG BASE</u>	<u>A200465 010</u>	Nov 29, 2013
AB		<u>EQ 10MG BASE;EQ 80MG BASE</u>	<u>A200465 011</u>	Nov 29, 2013
AB	ZYDUS PHARMS	<u>EQ 2.5MG BASE;EQ 10MG BASE</u>	<u>A207762 001</u>	Jan 11, 2019
AB		<u>EQ 2.5MG BASE;EQ 20MG BASE</u>	<u>A207762 002</u>	Jan 11, 2019
AB		<u>EQ 2.5MG BASE;EQ 40MG BASE</u>	<u>A207762 003</u>	Jan 11, 2019
AB		<u>EQ 5MG BASE;EQ 10MG BASE</u>	<u>A207762 004</u>	Jan 11, 2019
AB		<u>EQ 5MG BASE;EQ 20MG BASE</u>	<u>A207762 005</u>	Jan 11, 2019
AB		<u>EQ 5MG BASE;EQ 40MG BASE</u>	<u>A207762 006</u>	Jan 11, 2019
AB		<u>EQ 5MG BASE;EQ 80MG BASE</u>	<u>A207762 007</u>	Jan 11, 2019
AB		<u>EQ 10MG BASE;EQ 10MG BASE</u>	<u>A207762 008</u>	Jan 11, 2019
AB		<u>EQ 10MG BASE;EQ 20MG BASE</u>	<u>A207762 009</u>	Jan 11, 2019
AB		<u>EQ 10MG BASE;EQ 40MG BASE</u>	<u>A207762 010</u>	Jan 11, 2019
AB		<u>EQ 10MG BASE;EQ 80MG BASE</u>	<u>A207762 011</u>	Jan 11, 2019

CADUET

AB	+	PHARMACIA	<u>EQ 2.5MG BASE;EQ 10MG BASE</u>	<u>N021540 009</u>	Jul 29, 2004
AB	+		<u>EQ 2.5MG BASE;EQ 20MG BASE</u>	<u>N021540 010</u>	Jul 29, 2004
AB	+		<u>EQ 2.5MG BASE;EQ 40MG BASE</u>	<u>N021540 011</u>	Jul 29, 2004
AB	+		<u>EQ 5MG BASE;EQ 10MG BASE</u>	<u>N021540 001</u>	Jan 30, 2004
AB	+		<u>EQ 5MG BASE;EQ 20MG BASE</u>	<u>N021540 002</u>	Jan 30, 2004
AB	+		<u>EQ 5MG BASE;EQ 40MG BASE</u>	<u>N021540 003</u>	Jan 30, 2004
AB	+		<u>EQ 5MG BASE;EQ 80MG BASE</u>	<u>N021540 004</u>	Jan 30, 2004
AB	+		<u>EQ 10MG BASE;EQ 10MG BASE</u>	<u>N021540 005</u>	Jan 30, 2004
AB	+		<u>EQ 10MG BASE;EQ 20MG BASE</u>	<u>N021540 006</u>	Jan 30, 2004
AB	+		<u>EQ 10MG BASE;EQ 40MG BASE</u>	<u>N021540 007</u>	Jan 30, 2004
AB	+		<u>EQ 10MG BASE;EQ 80MG BASE</u>	<u>N021540 008</u>	Jan 30, 2004

AMLODIPINE BESYLATE; BENAZEPRIL HYDROCHLORIDE

CAPSULE; ORAL

AMLODIPINE BESYLATE AND BENAZEPRIL HYDROCHLORIDE

AB	AUROBINDO PHARMA LTD	<u>EQ 2.5MG BASE;10MG</u>	<u>A202239 001</u>	Sep 05, 2012
AB		<u>EQ 5MG BASE;10MG</u>	<u>A202239 002</u>	Sep 05, 2012
AB		<u>EQ 5MG BASE;20MG</u>	<u>A202239 003</u>	Sep 05, 2012
AB		<u>EQ 5MG BASE;40MG</u>	<u>A202239 004</u>	Sep 05, 2012
AB		<u>EQ 10MG BASE;20MG</u>	<u>A202239 005</u>	Sep 05, 2012
AB		<u>EQ 10MG BASE;40MG</u>	<u>A202239 006</u>	Sep 05, 2012
AB	DR REDDYS LABS INC	<u>EQ 2.5MG BASE;10MG</u>	<u>A077183 001</u>	Apr 15, 2010
AB		<u>EQ 5MG BASE;10MG</u>	<u>A077183 002</u>	Apr 15, 2010
AB		<u>EQ 5MG BASE;20MG</u>	<u>A077183 003</u>	Apr 15, 2010
AB		<u>EQ 5MG BASE;40MG</u>	<u>A090149 001</u>	Jul 05, 2011
AB		<u>EQ 10MG BASE;20MG</u>	<u>A077183 004</u>	Apr 15, 2010
AB		<u>EQ 10MG BASE;40MG</u>	<u>A090149 002</u>	Jul 05, 2011
AB	HERITAGE	<u>EQ 2.5MG BASE;10MG</u>	<u>A091431 001</u>	Dec 30, 2013
AB		<u>EQ 5MG BASE;10MG</u>	<u>A091431 002</u>	Dec 30, 2013
AB		<u>EQ 5MG BASE;20MG</u>	<u>A091431 003</u>	Dec 30, 2013
AB		<u>EQ 5MG BASE;40MG</u>	<u>A091431 004</u>	Dec 30, 2013
AB		<u>EQ 10MG BASE;20MG</u>	<u>A091431 005</u>	Dec 30, 2013
AB		<u>EQ 10MG BASE;40MG</u>	<u>A091431 006</u>	Dec 30, 2013
AB	LUPIN PHARMS	<u>EQ 2.5MG BASE;10MG</u>	<u>A078466 001</u>	Feb 05, 2010
AB		<u>EQ 5MG BASE;10MG</u>	<u>A078466 002</u>	Feb 05, 2010
AB		<u>EQ 5MG BASE;20MG</u>	<u>A078466 003</u>	Feb 05, 2010
AB		<u>EQ 5MG BASE;40MG</u>	<u>A078466 005</u>	Jul 05, 2011
AB		<u>EQ 10MG BASE;20MG</u>	<u>A078466 004</u>	Feb 05, 2010
AB		<u>EQ 10MG BASE;40MG</u>	<u>A078466 006</u>	Jul 05, 2011
AB	WATSON LABS	<u>EQ 2.5MG BASE;10MG</u>	<u>A077890 001</u>	Oct 14, 2010
AB		<u>EQ 5MG BASE;10MG</u>	<u>A077890 002</u>	Oct 14, 2010
AB		<u>EQ 5MG BASE;20MG</u>	<u>A077890 003</u>	Oct 14, 2010
AB		<u>EQ 10MG BASE;20MG</u>	<u>A077890 004</u>	Oct 14, 2010

PRESCRIPTION DRUG PRODUCT LIST

AMLODIPINE BESYLATE; BENAZEPRIL HYDROCHLORIDE

CAPSULE;ORAL

AMLODIPINE BESYLATE AND BENAZEPRIL HYDROCHLORIDE

<u>AB</u>	WATSON LABS INC	<u>EQ 5MG BASE;40MG</u>	<u>A090364 001</u>	Jul 05, 2011
<u>AB</u>		<u>EQ 10MG BASE;40MG</u>	<u>A090364 002</u>	Jul 05, 2011
<u>LOTREL</u>				
<u>AB</u>	+ SANDOZ	<u>EQ 2.5MG BASE;10MG</u>	<u>N020364 002</u>	Mar 03, 1995
<u>AB</u>	+	<u>EQ 5MG BASE;10MG</u>	<u>N020364 003</u>	Mar 03, 1995
<u>AB</u>	+	<u>EQ 5MG BASE;20MG</u>	<u>N020364 004</u>	Mar 03, 1995
<u>AB</u>	+	<u>EQ 5MG BASE;40MG</u>	<u>N020364 007</u>	Apr 11, 2006
<u>AB</u>	+	<u>EQ 10MG BASE;20MG</u>	<u>N020364 005</u>	Jun 20, 2002
<u>AB</u>	+	<u>EQ 10MG BASE;40MG</u>	<u>N020364 006</u>	Apr 11, 2006

AMLODIPINE BESYLATE; HYDROCHLOROTHIAZIDE; OLMESARTAN MEDOXOMIL

TABLET;ORAL

OLMESARTAN MEDOXOMIL, AMLODIPINE AND HYDROCHLOROTHIAZIDE

<u>AB</u>	ENDO OPERATIONS	<u>EQ 5MG BASE;12.5MG;20MG</u>	<u>A206137 001</u>	Oct 26, 2016
<u>AB</u>		<u>EQ 5MG BASE;12.5MG;40MG</u>	<u>A206137 002</u>	Oct 26, 2016
<u>AB</u>		<u>EQ 5MG BASE;25MG;40MG</u>	<u>A206137 003</u>	Oct 26, 2016
<u>AB</u>		<u>EQ 10MG BASE;12.5MG;40MG</u>	<u>A206137 004</u>	Oct 26, 2016
<u>AB</u>		<u>EQ 10MG BASE;25MG;40MG</u>	<u>A206137 005</u>	Oct 26, 2016
<u>AB</u>	TEVA PHARMS USA	<u>EQ 5MG BASE;12.5MG;20MG</u>	<u>A202491 001</u>	Nov 03, 2016
<u>AB</u>		<u>EQ 5MG BASE;12.5MG;40MG</u>	<u>A202491 002</u>	Nov 03, 2016
<u>AB</u>		<u>EQ 5MG BASE;25MG;40MG</u>	<u>A202491 003</u>	Nov 03, 2016
<u>AB</u>		<u>EQ 10MG BASE;12.5MG;40MG</u>	<u>A202491 004</u>	Nov 03, 2016
<u>AB</u>		<u>EQ 10MG BASE;25MG;40MG</u>	<u>A202491 005</u>	Nov 03, 2016
<u>AB</u>	TORRENT	<u>EQ 5MG BASE;12.5MG;20MG</u>	<u>A203580 001</u>	Oct 26, 2016
<u>AB</u>		<u>EQ 5MG BASE;12.5MG;40MG</u>	<u>A203580 002</u>	Oct 26, 2016
<u>AB</u>		<u>EQ 5MG BASE;25MG;40MG</u>	<u>A203580 003</u>	Oct 26, 2016
<u>AB</u>		<u>EQ 10MG BASE;12.5MG;40MG</u>	<u>A203580 004</u>	Oct 26, 2016
<u>AB</u>		<u>EQ 10MG BASE;25MG;40MG</u>	<u>A203580 005</u>	Oct 26, 2016
<u>TRIBENZOR</u>				
<u>AB</u>	+ COSETTE	<u>EQ 5MG BASE;12.5MG;20MG</u>	<u>N200175 001</u>	Jul 23, 2010
<u>AB</u>	+	<u>EQ 5MG BASE;12.5MG;40MG</u>	<u>N200175 002</u>	Jul 23, 2010
<u>AB</u>	+	<u>EQ 5MG BASE;25MG;40MG</u>	<u>N200175 003</u>	Jul 23, 2010
<u>AB</u>	+	<u>EQ 10MG BASE;12.5MG;40MG</u>	<u>N200175 004</u>	Jul 23, 2010
<u>AB</u>	+	<u>EQ 10MG BASE;25MG;40MG</u>	<u>N200175 005</u>	Jul 23, 2010

AMLODIPINE BESYLATE; HYDROCHLOROTHIAZIDE; VALSARTAN

TABLET;ORAL

AMLODIPINE BESYLATE, VALSARTAN AND HYDROCHLOROTHIAZIDE

<u>AB</u>	AUROBINDO PHARMA	<u>EQ 5MG BASE;12.5MG;160MG</u>	<u>A206180 001</u>	Dec 19, 2017
<u>AB</u>		<u>EQ 5MG BASE;25MG;160MG</u>	<u>A206180 002</u>	Dec 19, 2017
<u>AB</u>		<u>EQ 10MG BASE;12.5MG;160MG</u>	<u>A206180 003</u>	Dec 19, 2017
<u>AB</u>		<u>EQ 10MG BASE;25MG;160MG</u>	<u>A206180 004</u>	Dec 19, 2017
<u>AB</u>		<u>EQ 10MG BASE;25MG;320MG</u>	<u>A206180 005</u>	Dec 19, 2017
<u>AB</u>	LUPIN LTD	<u>EQ 5MG BASE;12.5MG;160MG</u>	<u>A200797 001</u>	Jun 03, 2015
<u>AB</u>		<u>EQ 5MG BASE;25MG;160MG</u>	<u>A200797 002</u>	Jun 03, 2015
<u>AB</u>		<u>EQ 10MG BASE;12.5MG;160MG</u>	<u>A200797 003</u>	Jun 03, 2015
<u>AB</u>		<u>EQ 10MG BASE;25MG;160MG</u>	<u>A200797 004</u>	Jun 03, 2015
<u>AB</u>		<u>EQ 10MG BASE;25MG;320MG</u>	<u>A200797 005</u>	Jun 03, 2015
<u>AB</u>	MACLEODS PHARMS LTD	<u>EQ 5MG BASE;12.5MG;160MG</u>	<u>A207299 001</u>	Jan 07, 2025
<u>AB</u>		<u>EQ 5MG BASE;25MG;160MG</u>	<u>A207299 002</u>	Jan 07, 2025
<u>AB</u>		<u>EQ 10MG BASE;12.5MG;160MG</u>	<u>A207299 003</u>	Jan 07, 2025
<u>AB</u>		<u>EQ 10MG BASE;25MG;160MG</u>	<u>A207299 004</u>	Jan 07, 2025
<u>AB</u>		<u>EQ 10MG BASE;25MG;320MG</u>	<u>A207299 005</u>	Jan 07, 2025
<u>AB</u>	STRIDES PHARMA	<u>EQ 5MG BASE;12.5MG;160MG</u>	<u>A201087 001</u>	Jun 01, 2015
<u>AB</u>		<u>EQ 5MG BASE;25MG;160MG</u>	<u>A201087 002</u>	Jun 01, 2015
<u>AB</u>		<u>EQ 10MG BASE;12.5MG;160MG</u>	<u>A201087 003</u>	Jun 01, 2015
<u>AB</u>		<u>EQ 10MG BASE;25MG;160MG</u>	<u>A201087 004</u>	Jun 01, 2015
<u>AB</u>		<u>EQ 10MG BASE;25MG;320MG</u>	<u>A201087 005</u>	Jun 01, 2015
<u>EXFORGE HCT</u>				
<u>AB</u>	+ NOVARTIS	<u>EQ 5MG BASE;12.5MG;160MG</u>	<u>N022314 001</u>	Apr 30, 2009
<u>AB</u>	+	<u>EQ 5MG BASE;25MG;160MG</u>	<u>N022314 002</u>	Apr 30, 2009
<u>AB</u>	+	<u>EQ 10MG BASE;12.5MG;160MG</u>	<u>N022314 003</u>	Apr 30, 2009
<u>AB</u>	+	<u>EQ 10MG BASE;25MG;160MG</u>	<u>N022314 004</u>	Apr 30, 2009
<u>AB</u>	+	<u>EQ 10MG BASE;25MG;320MG</u>	<u>N022314 005</u>	Apr 30, 2009

PRESCRIPTION DRUG PRODUCT LIST

AMLODIPINE BESYLATE; OLMESARTAN MEDOXOMIL

TABLET; ORAL

AMLODIPINE AND OLMESARTAN MEDOXOMIL

<u>AB</u>	AJANTA PHARMA LTD	<u>EQ 5MG BASE;20MG</u>	<u>A207216 001</u>	Oct 28, 2016
<u>AB</u>		<u>EQ 5MG BASE;40MG</u>	<u>A207216 002</u>	Oct 28, 2016
<u>AB</u>		<u>EQ 10MG BASE;20MG</u>	<u>A207216 003</u>	Oct 28, 2016
<u>AB</u>		<u>EQ 10MG BASE;40MG</u>	<u>A207216 004</u>	Oct 28, 2016
<u>AB</u>	ALEMBIC	<u>EQ 5MG BASE;20MG</u>	<u>A207073 001</u>	Jul 17, 2017
<u>AB</u>		<u>EQ 5MG BASE;40MG</u>	<u>A207073 002</u>	Jul 17, 2017
<u>AB</u>		<u>EQ 10MG BASE;20MG</u>	<u>A207073 003</u>	Jul 17, 2017
<u>AB</u>		<u>EQ 10MG BASE;40MG</u>	<u>A207073 004</u>	Jul 17, 2017
<u>AB</u>	ALKEM LABS LTD	<u>EQ 5MG BASE;20MG</u>	<u>A209042 001</u>	Aug 14, 2017
<u>AB</u>		<u>EQ 5MG BASE;40MG</u>	<u>A209042 002</u>	Aug 14, 2017
<u>AB</u>		<u>EQ 10MG BASE;20MG</u>	<u>A209042 003</u>	Aug 14, 2017
<u>AB</u>		<u>EQ 10MG BASE;40MG</u>	<u>A209042 004</u>	Aug 14, 2017
<u>AB</u>	AUROBINDO PHARMA	<u>EQ 5MG BASE;20MG</u>	<u>A206906 001</u>	May 15, 2017
<u>AB</u>		<u>EQ 5MG BASE;40MG</u>	<u>A206906 002</u>	May 15, 2017
<u>AB</u>		<u>EQ 10MG BASE;20MG</u>	<u>A206906 003</u>	May 15, 2017
<u>AB</u>		<u>EQ 10MG BASE;40MG</u>	<u>A206906 004</u>	May 15, 2017
<u>AB</u>	MACLEODS PHARMS LTD	<u>EQ 5MG BASE;20MG</u>	<u>A206884 001</u>	Oct 26, 2016
<u>AB</u>		<u>EQ 5MG BASE;40MG</u>	<u>A206884 003</u>	Oct 26, 2016
<u>AB</u>		<u>EQ 10MG BASE;20MG</u>	<u>A206884 002</u>	Oct 26, 2016
<u>AB</u>		<u>EQ 10MG BASE;40MG</u>	<u>A206884 004</u>	Oct 26, 2016
<u>AB</u>	MICRO LABS	<u>EQ 5MG BASE;20MG</u>	<u>A207435 001</u>	Nov 02, 2017
<u>AB</u>		<u>EQ 5MG BASE;40MG</u>	<u>A207435 002</u>	Nov 02, 2017
<u>AB</u>		<u>EQ 10MG BASE;20MG</u>	<u>A207435 003</u>	Nov 02, 2017
<u>AB</u>		<u>EQ 10MG BASE;40MG</u>	<u>A207435 004</u>	Nov 02, 2017
<u>AB</u>	SCIEGEN PHARMS INC	<u>EQ 5MG BASE;20MG</u>	<u>A209010 001</u>	Dec 03, 2018
<u>AB</u>		<u>EQ 5MG BASE;40MG</u>	<u>A209010 002</u>	Dec 03, 2018
<u>AB</u>		<u>EQ 10MG BASE;20MG</u>	<u>A209010 003</u>	Dec 03, 2018
<u>AB</u>		<u>EQ 10MG BASE;40MG</u>	<u>A209010 004</u>	Dec 03, 2018
<u>AB</u>	ZYDUS PHARMS	<u>EQ 5MG BASE;20MG</u>	<u>A207771 001</u>	Sep 22, 2017
<u>AB</u>		<u>EQ 5MG BASE;40MG</u>	<u>A207771 002</u>	Sep 22, 2017
<u>AB</u>		<u>EQ 10MG BASE;20MG</u>	<u>A207771 003</u>	Sep 22, 2017
<u>AB</u>		<u>EQ 10MG BASE;40MG</u>	<u>A207771 004</u>	Sep 22, 2017
<u>AZOR</u>				
<u>AB</u>	+ COSETTE	<u>EQ 5MG BASE;20MG</u>	<u>N022100 001</u>	Sep 26, 2007
<u>AB</u>	+	<u>EQ 5MG BASE;40MG</u>	<u>N022100 002</u>	Sep 26, 2007
<u>AB</u>	+	<u>EQ 10MG BASE;20MG</u>	<u>N022100 003</u>	Sep 26, 2007
<u>AB</u>	+!	<u>EQ 10MG BASE;40MG</u>	<u>N022100 004</u>	Sep 26, 2007

AMLODIPINE BESYLATE; TELMISARTAN

TABLET; ORAL

TELMISARTAN AND AMLODIPINE

<u>AB</u>	LUPIN LTD	<u>EQ 5MG BASE;40MG</u>	<u>A201586 001</u>	Jan 08, 2014
<u>AB</u>		<u>EQ 5MG BASE;80MG</u>	<u>A201586 003</u>	Jan 08, 2014
<u>AB</u>		<u>EQ 10MG BASE;40MG</u>	<u>A201586 002</u>	Jan 08, 2014
<u>AB</u>		<u>EQ 10MG BASE;80MG</u>	<u>A201586 004</u>	Jan 08, 2014
<u>AB</u>	MYLAN	<u>EQ 5MG BASE;40MG</u>	<u>A202516 001</u>	Aug 26, 2014
<u>AB</u>		<u>EQ 5MG BASE;80MG</u>	<u>A202516 003</u>	Aug 26, 2014
<u>AB</u>		<u>EQ 10MG BASE;40MG</u>	<u>A202516 002</u>	Aug 26, 2014
<u>AB</u>	!	<u>EQ 10MG BASE;80MG</u>	<u>A202516 004</u>	Aug 26, 2014

AMLODIPINE BESYLATE; VALSARTAN

TABLET; ORAL

AMLODIPINE BESYLATE AND VALSARTAN

<u>AB</u>	ALEMBIC	<u>EQ 5MG BASE;160MG</u>	<u>A202713 001</u>	Apr 03, 2015
<u>AB</u>		<u>EQ 5MG BASE;320MG</u>	<u>A202713 003</u>	Apr 03, 2015
<u>AB</u>		<u>EQ 10MG BASE;160MG</u>	<u>A202713 002</u>	Apr 03, 2015
<u>AB</u>		<u>EQ 10MG BASE;320MG</u>	<u>A202713 004</u>	Apr 03, 2015
<u>AB</u>	AUROBINDO PHARMA	<u>EQ 5MG BASE;160MG</u>	<u>A206512 001</u>	Apr 22, 2016
<u>AB</u>		<u>EQ 5MG BASE;320MG</u>	<u>A206512 002</u>	Apr 22, 2016
<u>AB</u>		<u>EQ 10MG BASE;160MG</u>	<u>A206512 003</u>	Apr 22, 2016
<u>AB</u>		<u>EQ 10MG BASE;320MG</u>	<u>A206512 004</u>	Apr 22, 2016
<u>AB</u>	HETERO LABS	<u>EQ 5MG BASE;160MG</u>	<u>A205137 001</u>	Sep 16, 2016
<u>AB</u>		<u>EQ 5MG BASE;320MG</u>	<u>A205137 003</u>	Sep 16, 2016
<u>AB</u>		<u>EQ 10MG BASE;160MG</u>	<u>A205137 002</u>	Sep 16, 2016
<u>AB</u>		<u>EQ 10MG BASE;320MG</u>	<u>A205137 004</u>	Sep 16, 2016
<u>AB</u>	LUPIN	<u>EQ 5MG BASE;160MG</u>	<u>A090245 001</u>	Mar 30, 2015
<u>AB</u>		<u>EQ 5MG BASE;320MG</u>	<u>A090245 003</u>	Mar 30, 2015
<u>AB</u>		<u>EQ 10MG BASE;160MG</u>	<u>A090245 002</u>	Mar 30, 2015
<u>AB</u>		<u>EQ 10MG BASE;320MG</u>	<u>A090245 004</u>	Mar 30, 2015
<u>AB</u>	MACLEODS PHARMS LTD	<u>EQ 5MG BASE;160MG</u>	<u>A207292 001</u>	Oct 11, 2024

PRESCRIPTION DRUG PRODUCT LIST

AMLODIPINE BESYLATE; VALSARTAN

TABLET; ORAL

AMLODIPINE BESYLATE AND VALSARTAN

<u>AB</u>		<u>EQ 5MG BASE;320MG</u>	<u>A207292 003</u>	Oct 11, 2024
<u>AB</u>		<u>EQ 10MG BASE;160MG</u>	<u>A207292 002</u>	Oct 11, 2024
<u>AB</u>		<u>EQ 10MG BASE;320MG</u>	<u>A207292 004</u>	Oct 11, 2024
<u>AB</u>	MYLAN	<u>EQ 5MG BASE;160MG</u>	<u>A090483 001</u>	Mar 30, 2015
<u>AB</u>		<u>EQ 5MG BASE;320MG</u>	<u>A090483 003</u>	Mar 30, 2015
<u>AB</u>		<u>EQ 10MG BASE;160MG</u>	<u>A090483 002</u>	Mar 30, 2015
<u>AB</u>		<u>EQ 10MG BASE;320MG</u>	<u>A090483 004</u>	Mar 30, 2015
<u>AB</u>	NOVEL LABS INC	<u>EQ 5MG BASE;160MG</u>	<u>A202829 001</u>	Mar 30, 2015
<u>AB</u>		<u>EQ 5MG BASE;320MG</u>	<u>A202829 003</u>	Mar 30, 2015
<u>AB</u>		<u>EQ 10MG BASE;160MG</u>	<u>A202829 002</u>	Mar 30, 2015
<u>AB</u>		<u>EQ 10MG BASE;320MG</u>	<u>A202829 004</u>	Mar 30, 2015
<u>AB</u>	STRIDES PHARMA	<u>EQ 5MG BASE;160MG</u>	<u>A090011 001</u>	Mar 28, 2013
<u>AB</u>		<u>EQ 5MG BASE;320MG</u>	<u>A090011 003</u>	Mar 28, 2013
<u>AB</u>		<u>EQ 10MG BASE;160MG</u>	<u>A090011 002</u>	Mar 28, 2013
<u>AB</u>		<u>EQ 10MG BASE;320MG</u>	<u>A090011 004</u>	Mar 28, 2013
<u>EXFORGE</u>				
<u>AB</u>	+ NOVARTIS	<u>EQ 5MG BASE;160MG</u>	<u>N021990 002</u>	Jun 20, 2007
<u>AB</u>	+	<u>EQ 5MG BASE;320MG</u>	<u>N021990 004</u>	Jun 20, 2007
<u>AB</u>	+!	<u>EQ 10MG BASE;160MG</u>	<u>N021990 003</u>	Jun 20, 2007
<u>AB</u>	+!	<u>EQ 10MG BASE;320MG</u>	<u>N021990 005</u>	Jun 20, 2007

AMMONIA N-13

INJECTABLE; INTRAVENOUS

AMMONIA N 13

<u>AP</u>	3D IMAGING DRUG	<u>30mCi-300mCi/8ML (3.75-37.5mCi/ML)</u>	<u>A203779 001</u>	Oct 19, 2015
<u>AP</u>	BIOMEDCL RES FDN	<u>48.75mCi-487.5mCi/13ML (3.75-37.5mCi/ML)</u>	<u>A204352 001</u>	May 01, 2015
<u>AP</u>	BRIGHAM WOMENS HOSP	<u>30mCi-300mCi/8ML (3.75-37.5mCi/ML)</u>	<u>A203783 001</u>	Oct 30, 2014
<u>AP</u>	CARDINAL HLTH 414	<u>30mCi-300mCi/8ML (3.75-37.5mCi/ML)</u>	<u>A203700 001</u>	Feb 25, 2013
<u>AP</u>	DECATUR	<u>30mCi-300mCi/8ML (3.75-37.5mCi/ML)</u>	<u>A204465 001</u>	Oct 23, 2014
<u>AP</u>	+! FEINSTEIN	<u>30mCi-300mCi/8ML (3.75-37.5mCi/ML)</u>	<u>N022119 001</u>	Aug 23, 2007
<u>AP</u>	GEN HOSP	<u>30mCi-300mCi/8ML (3.75-37.5mCi/ML)</u>	<u>A207025 001</u>	Feb 03, 2016
<u>AP</u>	IONETIX	<u>22.5mCi-225mCi/6ML (3.75-37.5mCi/ML)</u>	<u>A210524 001</u>	Dec 21, 2018
<u>AP</u>	JOHNS HOPKINS UNIV	<u>30mCi-300mCi/8ML (3.75-37.5mCi/ML)</u>	<u>A204514 001</u>	Aug 19, 2014
<u>AP</u>	KREITCHMAN PET CTR	<u>30mCi-300mCi/8ML (3.75-37.5mCi/ML)</u>	<u>A203938 001</u>	Dec 09, 2013
<u>AP</u>	MCPRF	<u>30mCi-300mCi/8ML (3.75-37.5mCi/ML)</u>	<u>A203321 001</u>	Feb 25, 2013
<u>AP</u>	METHODIST	<u>3.75mCi-260mCi/ML</u>	<u>A215083 001</u>	Jul 09, 2021
<u>AP</u>	MIDWEST MEDCL	<u>30mCi-300mCi/8ML (3.75-37.5mCi/ML)</u>	<u>A204457 001</u>	Nov 18, 2015
<u>AP</u>	MIPS CRF	<u>30mCi-300mCi/8ML (3.75-37.5mCi/ML)</u>	<u>A204535 001</u>	Nov 20, 2014
<u>AP</u>	NCM USA BRONX LLC	<u>3.75mCi-260mCi/ML</u>	<u>A204515 001</u>	Feb 04, 2015
<u>AP</u>	NUKEMED	<u>30mCi-300mCi/8ML (3.75-37.5mCi/ML)</u>	<u>A204455 001</u>	Apr 23, 2015
<u>AP</u>	PETNET	<u>30mCi-300mCi (3.75-37.5mCi/ML)</u>	<u>A204510 001</u>	Nov 02, 2015
<u>AP</u>	PRECISION NUCLEAR	<u>3.75mCi-260mCi/ML</u>	<u>A204547 001</u>	Aug 14, 2015
<u>AP</u>	! SOFIE	<u>3.75mCi-260mCi/ML</u>	<u>A203543 001</u>	Dec 14, 2012
<u>AP</u>	UCLA BIOMEDICAL	<u>30mCi-300mCi/8ML (3.75-37.5mCi/ML)</u>	<u>A203812 001</u>	Jun 27, 2013
<u>AP</u>	UCSF RODIOPHARM	<u>30mCi-300mCi/8ML (3.75-37.5mCi/ML)</u>	<u>A204496 001</u>	Mar 28, 2014
<u>AP</u>	UNIV ALAHAMA BIRM	<u>15mCi-150mCi/4ML (3.75-37.5mCi/ML)</u>	<u>A211698 001</u>	Nov 03, 2022
<u>AP</u>	UNIV TX SW MEDCTR	<u>30mCi-300mCi/8ML (3.75-37.5mCi/ML)</u>	<u>A209507 001</u>	Nov 01, 2019
<u>AP</u>	UNIV WISCONSIN	<u>30mCi-300mCi/8ML (3.75-37.5mCi/ML)</u>	<u>A211740 001</u>	Sep 09, 2020
<u>AP</u>	WA UNIV SCH MED	<u>30mCi-300mCi/8ML (3.75-37.5mCi/ML)</u>	<u>A204506 001</u>	Feb 07, 2014

AMMONIUM CHLORIDE

INJECTABLE; INJECTION

AMMONIUM CHLORIDE IN PLASTIC CONTAINER

! HOSPIRA

5MEQ/ML

A088366 001 Jun 13, 1984

AMMONIUM LACTATE

CREAM; TOPICAL

AMMONIUM LACTATE

<u>AB</u>	! PADAGIS ISRAEL	<u>EQ 12% BASE</u>	<u>A075774 001</u>	May 01, 2002
<u>AB</u>	TARO	<u>EQ 12% BASE</u>	<u>A075883 001</u>	Apr 10, 2003
LOTION; TOPICAL				
<u>AMMONIUM LACTATE</u>				
<u>AB</u>	! PADAGIS ISRAEL	<u>EQ 12% BASE</u>	<u>A075570 001</u>	Jun 23, 2004
<u>AB</u>	TARO	<u>EQ 12% BASE</u>	<u>A076216 001</u>	May 28, 2004

PRESCRIPTION DRUG PRODUCT LIST

AMOXAPINE

TABLET; ORAL

AMOXAPINE

<u>AB</u>	CHARTWELL RX	<u>25MG</u>	<u>A072879 002</u>	Jun 28, 1991
<u>AB</u>		<u>50MG</u>	<u>A072879 003</u>	Jun 28, 1991
<u>AB</u>		<u>100MG</u>	<u>A072879 004</u>	Jun 28, 1991
<u>AB</u>		<u>150MG</u>	<u>A072879 001</u>	Jun 28, 1991
<u>AB</u>	WATSON LABS	<u>25MG</u>	<u>A072691 002</u>	Aug 28, 1992
<u>AB</u>		<u>50MG</u>	<u>A072691 003</u>	Aug 28, 1992
<u>AB</u>		<u>100MG</u>	<u>A072691 004</u>	Aug 28, 1992
<u>AB</u>	!	<u>150MG</u>	<u>A072691 001</u>	Aug 28, 1992

AMOXICILLIN

CAPSULE; ORAL

AMOXICILLIN

<u>AB</u>	AUROBINDO	<u>250MG</u>	<u>A065271 001</u>	Nov 09, 2005
<u>AB</u>		<u>500MG</u>	<u>A065271 002</u>	Nov 09, 2005
<u>AB</u>	CHARTWELL	<u>250MG</u>	<u>A062058 001</u>	
<u>AB</u>		<u>500MG</u>	<u>A062058 002</u>	
<u>AB</u>	HIKMA PHARMS	<u>250MG</u>	<u>A065291 001</u>	Feb 05, 2007
<u>AB</u>		<u>500MG</u>	<u>A065291 002</u>	Feb 05, 2007
<u>AB</u>	MICRO LABS	<u>250MG</u>	<u>A207471 001</u>	Jun 24, 2022
<u>AB</u>		<u>500MG</u>	<u>A207471 002</u>	Jun 24, 2022
<u>AB</u>	SANDOZ	<u>250MG</u>	<u>A064076 001</u>	Sep 30, 1994
<u>AB</u>		<u>500MG</u>	<u>A064076 002</u>	Sep 30, 1994
<u>AB</u>	TEVA	<u>250MG</u>	<u>A061926 001</u>	
<u>AB</u>	!	<u>500MG</u>	<u>A061926 003</u>	

AMOXIL

<u>AB</u>	US ANTIBIOTICS	<u>250MG</u>	<u>A062216 001</u>	
<u>AB</u>		<u>500MG</u>	<u>A062216 004</u>	

FOR SUSPENSION; ORAL

AMOXICILLIN

<u>AB</u>	AUROBINDO	<u>200MG/5ML</u>	<u>A065334 001</u>	Dec 28, 2006
<u>AB</u>		<u>400MG/5ML</u>	<u>A065334 002</u>	Dec 28, 2006
<u>AB</u>	AUROBINDO PHARMA LTD	<u>125MG/5ML</u>	<u>A204030 001</u>	Sep 15, 2014
<u>AB</u>		<u>250MG/5ML</u>	<u>A204030 002</u>	Sep 15, 2014
<u>AB</u>	CHARTWELL RX	<u>400MG/5ML</u>	<u>A065319 002</u>	Jun 18, 2007
<u>AB</u>	HIKMA	<u>125MG/5ML</u>	<u>A065322 002</u>	Jun 19, 2006
<u>AB</u>		<u>200MG/5ML</u>	<u>A065325 002</u>	Jun 19, 2006
<u>AB</u>		<u>250MG/5ML</u>	<u>A065322 001</u>	Jun 19, 2006
<u>AB</u>		<u>400MG/5ML</u>	<u>A065325 001</u>	Jun 19, 2006
<u>AB</u>	SANDOZ	<u>125MG/5ML</u>	<u>A065387 001</u>	Mar 26, 2007
<u>AB</u>		<u>200MG/5ML</u>	<u>A065378 001</u>	Mar 26, 2007
<u>AB</u>		<u>250MG/5ML</u>	<u>A065387 002</u>	Mar 26, 2007
<u>AB</u>		<u>400MG/5ML</u>	<u>A065378 002</u>	Mar 26, 2007
<u>AB</u>	TEVA	<u>200MG/5ML</u>	<u>A065119 001</u>	Dec 04, 2002
<u>AB</u>	!	<u>250MG/5ML</u>	<u>A061931 002</u>	
<u>AB</u>	!	<u>400MG/5ML</u>	<u>A065119 002</u>	Dec 04, 2002

AMOXICILLIN PEDIATRIC

<u>AB</u>	TEVA	<u>50MG/ML</u>	<u>A061931 003</u>	Dec 01, 1982
-----------	------	----------------	--------------------	--------------

AMOXIL

<u>AB</u>	US ANTIBIOTICS	<u>50MG/ML</u>	<u>A062226 005</u>	
<u>AB</u>		<u>125MG/5ML</u>	<u>A062226 001</u>	
<u>AB</u>		<u>250MG/5ML</u>	<u>A062226 002</u>	
<u>AB</u>	+	<u>400MG/5ML</u>	<u>N050760 002</u>	Apr 15, 1999

LAROTID

<u>AB</u>	US ANTIBIOTICS	<u>125MG/5ML</u>	<u>A062226 003</u>	
<u>AB</u>		<u>250MG/5ML</u>	<u>A062226 004</u>	

TABLET; ORAL

AMOXICILLIN

<u>AB</u>	AUROBINDO	<u>500MG</u>	<u>A065256 001</u>	Nov 09, 2005
<u>AB</u>		<u>875MG</u>	<u>A065256 002</u>	Nov 09, 2005
<u>AB</u>	HIKMA	<u>875MG</u>	<u>A065255 001</u>	Mar 29, 2006
<u>AB</u>	SANDOZ	<u>500MG</u>	<u>A065228 001</u>	Jul 13, 2005
<u>AB</u>		<u>875MG</u>	<u>A065228 002</u>	Jul 13, 2005
<u>AB</u>	TEVA	<u>500MG</u>	<u>A065056 001</u>	Sep 18, 2000
<u>AB</u>	!	<u>875MG</u>	<u>A065056 002</u>	Sep 18, 2000

TABLET, CHEWABLE; ORAL

AMOXICILLIN

	TEVA	125MG	A064013 002	Sep 11, 1995
	!	250MG	A064013 001	Dec 22, 1992

PRESCRIPTION DRUG PRODUCT LIST

AMOXICILLIN; CLARITHROMYCIN; LANSOPRAZOLE

CAPSULE, TABLET, CAPSULE, DELAYED REL PELLETS;ORAL

LANSOPRAZOLE, AMOXICILLIN AND CLARITHROMYCIN (COPACKAGED)

<u>AB</u>	!	RISING	<u>500MG;500MG;30MG</u>	<u>A206006</u>	<u>001</u>	Oct 07, 2016
<u>AB</u>		SANDOZ	<u>500MG;500MG;30MG</u>	<u>A202588</u>	<u>001</u>	Mar 04, 2014

AMOXICILLIN; CLARITHROMYCIN; VONOPRAZAN FUMARATE

CAPSULE, TABLET, TABLET;ORAL

VOQUEZNA TRIPLE PAK

+	!	PHATHOM	500MG;500MG;EQ 20MG BASE	N215152	001	May 03, 2022
---	---	---------	--------------------------	---------	-----	--------------

AMOXICILLIN; CLAVULANATE POTASSIUM

FOR SUSPENSION;ORAL

AMOXICILLIN AND CLAVULANATE POTASSIUM

<u>AB</u>		AUROBINDO PHARMA	<u>125MG/5ML;EQ 31.25MG BASE/5ML</u>	<u>A209371</u>	<u>001</u>	Apr 19, 2019
<u>AB</u>			<u>250MG/5ML;EQ 62.5MG BASE/5ML</u>	<u>A209371</u>	<u>002</u>	Apr 19, 2019
<u>AB</u>		AUROBINDO PHARMA LTD	<u>200MG/5ML;EQ 28.5MG BASE/5ML</u>	<u>A201090</u>	<u>001</u>	Dec 20, 2011
<u>AB</u>			<u>400MG/5ML;EQ 57MG BASE/5ML</u>	<u>A201090</u>	<u>002</u>	Dec 20, 2011
<u>AB</u>			<u>600MG/5ML;EQ 42.9MG BASE/5ML</u>	<u>A201091</u>	<u>001</u>	Dec 20, 2011
<u>AB</u>		CIPLA	<u>250MG/5ML;EQ 62.5MG BASE/5ML</u>	<u>A065431</u>	<u>001</u>	Nov 25, 2008
<u>AB</u>			<u>600MG/5ML;EQ 42.9MG BASE/5ML</u>	<u>A065420</u>	<u>001</u>	Dec 02, 2013
<u>AB</u>		DEVA HOLDING AS	<u>250MG/5ML;EQ 62.5MG BASE/5ML</u>	<u>A210374</u>	<u>001</u>	Nov 29, 2023
<u>AB</u>			<u>400MG/5ML;EQ 57MG BASE/5ML</u>	<u>A210416</u>	<u>001</u>	Dec 11, 2023
<u>AB</u>			<u>600MG/5ML;EQ 42.9MG BASE/5ML</u>	<u>A209351</u>	<u>001</u>	Sep 13, 2023
<u>AB</u>		HIKMA PHARMS	<u>200MG/5ML;EQ 28.5MG BASE/5ML</u>	<u>A065191</u>	<u>002</u>	Jan 25, 2005
<u>AB</u>			<u>400MG/5ML;EQ 57MG BASE/5ML</u>	<u>A065191</u>	<u>001</u>	Jan 25, 2005
<u>AB</u>			<u>600MG/5ML;EQ 42.9MG BASE/5ML</u>	<u>A065373</u>	<u>001</u>	Nov 09, 2007
<u>AB</u>		MICRO LABS	<u>600MG/5ML;EQ 42.9MG BASE/5ML</u>	<u>A217805</u>	<u>001</u>	Dec 08, 2023
<u>AB</u>		MICRO LABS LTD INDIA	<u>200MG/5ML;EQ 28.5MG BASE/5ML</u>	<u>A205187</u>	<u>001</u>	May 21, 2021
<u>AB</u>			<u>400MG/5ML;EQ 57MG BASE/5ML</u>	<u>A205187</u>	<u>002</u>	May 21, 2021
<u>AB</u>		SANDOZ	<u>200MG/5ML;EQ 28.5MG BASE/5ML</u>	<u>A065066</u>	<u>001</u>	Jun 05, 2002
<u>AB</u>			<u>400MG/5ML;EQ 57MG BASE/5ML</u>	<u>A065066</u>	<u>002</u>	Jun 05, 2002
<u>AB</u>		SANDOZ INC	<u>200MG/5ML;EQ 28.5MG BASE/5ML</u>	<u>A065098</u>	<u>001</u>	Dec 16, 2002
<u>AB</u>			<u>400MG/5ML;EQ 57MG BASE/5ML</u>	<u>A065098</u>	<u>002</u>	Dec 16, 2002
<u>AB</u>			<u>600MG/5ML;EQ 42.9MG BASE/5ML</u>	<u>A065358</u>	<u>001</u>	Aug 13, 2007
<u>AB</u>		TEVA	<u>200MG/5ML;EQ 28.5MG BASE/5ML</u>	<u>A065089</u>	<u>001</u>	May 25, 2004
<u>AB</u>	!		<u>400MG/5ML;EQ 57MG BASE/5ML</u>	<u>A065089</u>	<u>002</u>	May 25, 2004
<u>AB</u>	!		<u>600MG/5ML;EQ 42.9MG BASE/5ML</u>	<u>A065162</u>	<u>001</u>	Mar 12, 2004
<u>AUGMENTIN '125'</u>						
<u>AB</u>	+	US ANTIBIOTICS	<u>125MG/5ML;EQ 31.25MG BASE/5ML</u>	<u>N050575</u>	<u>001</u>	Aug 06, 1984
<u>AUGMENTIN '250'</u>						
<u>AB</u>	+	!	US ANTIBIOTICS	<u>250MG/5ML;EQ 62.5MG BASE/5ML</u>	<u>N050575</u>	<u>002</u> Aug 06, 1984
<u>AUGMENTIN ES-600</u>						
<u>AB</u>	+	US ANTIBIOTICS	<u>600MG/5ML;EQ 42.9MG BASE/5ML</u>	<u>N050755</u>	<u>001</u>	Jun 22, 2001
TABLET;ORAL						
<u>AMOXICILLIN AND CLAVULANATE POTASSIUM</u>						
<u>AB</u>		AUROBINDO PHARMA LTD	<u>250MG;EQ 125MG BASE</u>	<u>A091569</u>	<u>001</u>	Jan 20, 2012
<u>AB</u>	!		<u>500MG;EQ 125MG BASE</u>	<u>A091569</u>	<u>002</u>	Jan 20, 2012
<u>AB</u>			<u>875MG;EQ 125MG BASE</u>	<u>A091568</u>	<u>001</u>	Jan 20, 2012
<u>AB</u>		DEVA HOLDING AS	<u>500MG;EQ 125MG BASE</u>	<u>A209992</u>	<u>001</u>	Sep 15, 2023
<u>AB</u>			<u>875MG;EQ 125MG BASE</u>	<u>A209991</u>	<u>001</u>	Sep 15, 2023
<u>AB</u>		HIKMA PHARMS	<u>875MG;EQ 125MG BASE</u>	<u>A203824</u>	<u>001</u>	Aug 23, 2016
<u>AB</u>		MICRO LABS LTD INDIA	<u>250MG;EQ 125MG BASE</u>	<u>A205707</u>	<u>001</u>	Dec 30, 2016
<u>AB</u>			<u>500MG;EQ 125MG BASE</u>	<u>A205707</u>	<u>002</u>	Dec 30, 2016
<u>AB</u>			<u>875MG;EQ 125MG BASE</u>	<u>A204755</u>	<u>003</u>	Dec 30, 2016
<u>AB</u>	!	SANDOZ	<u>250MG;EQ 125MG BASE</u>	<u>A065189</u>	<u>001</u>	Aug 23, 2005
<u>AB</u>			<u>500MG;EQ 125MG BASE</u>	<u>A065064</u>	<u>001</u>	Mar 15, 2002
<u>AB</u>	!		<u>875MG;EQ 125MG BASE</u>	<u>A065063</u>	<u>001</u>	Mar 14, 2002
<u>AB</u>		SANDOZ INC	<u>875MG;EQ 125MG BASE</u>	<u>A065093</u>	<u>001</u>	Nov 21, 2002
<u>AB</u>		TEVA	<u>500MG;EQ 125MG BASE</u>	<u>A065101</u>	<u>001</u>	Oct 30, 2002
<u>AB</u>		TEVA PHARMS USA	<u>875MG;EQ 125MG BASE</u>	<u>A065096</u>	<u>001</u>	Oct 29, 2002
<u>AUGMENTIN '875'</u>						
<u>AB</u>	+	US ANTIBIOTICS	<u>875MG;EQ 125MG BASE</u>	<u>N050720</u>	<u>001</u>	Feb 13, 1996
TABLET, CHEWABLE;ORAL						
AMOXICILLIN AND CLAVULANATE POTASSIUM						
		TEVA	200MG;EQ 28.5MG BASE	A065205	001	Feb 09, 2005
	!		400MG;EQ 57MG BASE	A065205	002	Feb 09, 2005
TABLET, EXTENDED RELEASE;ORAL						
AMOXICILLIN AND CLAVULANATE POTASSIUM						
	!	SANDOZ	1GM;EQ 62.5MG BASE	A090227	001	Apr 21, 2010

PRESCRIPTION DRUG PRODUCT LIST

AMOXICILLIN; OMEPRAZOLE MAGNESIUM; RIFABUTIN

CAPSULE, DELAYED RELEASE;ORAL

TALICIA

+! REDHILL 250MG;EQ 10MG BASE;12.5MG N213004 001 Nov 01, 2019

AMOXICILLIN; VONOPRAZAN FUMARATE

CAPSULE, TABLET;ORAL

VOQUEZNA DUAL PAK

+! PHATHOM 500MG;EQ 20MG BASE N215153 001 May 03, 2022

AMPHETAMINE

TABLET, ORALLY DISINTEGRATING, EXTENDED RELEASE;ORAL

ADZENYS XR-ODT

+ NEOS THERAPS EQ 3.1MG BASE N204326 001 Jan 27, 2016
 + EQ 6.3MG BASE N204326 002 Jan 27, 2016
 + EQ 9.4MG BASE N204326 003 Jan 27, 2016
 + EQ 12.5MG BASE N204326 004 Jan 27, 2016
 + EQ 15.7MG BASE N204326 005 Jan 27, 2016
 +! EQ 18.8MG BASE N204326 006 Jan 27, 2016

AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATE

CAPSULE, EXTENDED RELEASE;ORAL

ADDERALL XR 10

AB1 + TAKEDA PHARMS USA **2.5MG;2.5MG;2.5MG;2.5MG** **N021303 001** Oct 11, 2001

ADDERALL XR 15

AB1 + TAKEDA PHARMS USA **3.75MG;3.75MG;3.75MG;3.75MG** **N021303 006** May 22, 2002

ADDERALL XR 20

AB1 + TAKEDA PHARMS USA **5MG;5MG;5MG;5MG** **N021303 002** Oct 11, 2001

ADDERALL XR 25

AB1 + TAKEDA PHARMS USA **6.25MG;6.25MG;6.25MG;6.25MG** **N021303 004** May 22, 2002

ADDERALL XR 30

AB1 +! TAKEDA PHARMS USA **7.5MG;7.5MG;7.5MG;7.5MG** **N021303 003** Oct 11, 2001

ADDERALL XR 5

AB1 + TAKEDA PHARMS USA **1.25MG;1.25MG;1.25MG;1.25MG** **N021303 005** May 22, 2002

DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE

AB1 ACTAVIS ELIZABETH **1.25MG;1.25MG;1.25MG;1.25MG** **A077302 001** Jun 22, 2012

AB1 **2.5MG;2.5MG;2.5MG;2.5MG** **A077302 002** Jun 22, 2012

AB1 **3.75MG;3.75MG;3.75MG;3.75MG** **A077302 003** Jun 22, 2012

AB1 **5MG;5MG;5MG;5MG** **A077302 004** Jun 22, 2012

AB1 **6.25MG;6.25MG;6.25MG;6.25MG** **A077302 005** Jun 22, 2012

AB1 **7.5MG;7.5MG;7.5MG;7.5MG** **A077302 006** Jun 22, 2012

AB1 ANI PHARMS **1.25MG;1.25MG;1.25MG;1.25MG** **A205401 001** Jan 22, 2019

AB1 **2.5MG;2.5MG;2.5MG;2.5MG** **A205401 002** Jan 22, 2019

AB1 **3.75MG;3.75MG;3.75MG;3.75MG** **A205401 003** Jan 22, 2019

AB1 **5MG;5MG;5MG;5MG** **A205401 004** Jan 22, 2019

AB1 **6.25MG;6.25MG;6.25MG;6.25MG** **A205401 005** Jan 22, 2019

AB1 **7.5MG;7.5MG;7.5MG;7.5MG** **A205401 006** Jan 22, 2019

AB1 ASCENT PHARMS INC **1.25MG;1.25MG;1.25MG;1.25MG** **A214959 001** Sep 29, 2021

AB1 **2.5MG;2.5MG;2.5MG;2.5MG** **A214959 002** Sep 29, 2021

AB1 **3.75MG;3.75MG;3.75MG;3.75MG** **A214959 003** Sep 29, 2021

AB1 **5MG;5MG;5MG;5MG** **A214959 004** Sep 29, 2021

AB1 **6.25MG;6.25MG;6.25MG;6.25MG** **A214959 005** Sep 29, 2021

AB1 **7.5MG;7.5MG;7.5MG;7.5MG** **A214959 006** Sep 29, 2021

AB1 ELITE LABS INC **1.25MG;1.25MG;1.25MG;1.25MG** **A212037 001** Dec 11, 2019

AB1 **2.5MG;2.5MG;2.5MG;2.5MG** **A212037 002** Dec 11, 2019

AB1 **3.75MG;3.75MG;3.75MG;3.75MG** **A212037 003** Dec 11, 2019

AB1 **5MG;5MG;5MG;5MG** **A212037 004** Dec 11, 2019

AB1 **6.25MG;6.25MG;6.25MG;6.25MG** **A212037 005** Dec 11, 2019

AB1 **7.5MG;7.5MG;7.5MG;7.5MG** **A212037 006** Dec 11, 2019

AB1 GRANULES **1.25MG;1.25MG;1.25MG;1.25MG** **A217027 001** Jan 23, 2023

AB1 **2.5MG;2.5MG;2.5MG;2.5MG** **A217027 002** Jan 23, 2023

AB1 **3.75MG;3.75MG;3.75MG;3.75MG** **A217027 003** Jan 23, 2023

AB1 **5MG;5MG;5MG;5MG** **A217027 004** Jan 23, 2023

AB1 **6.25MG;6.25MG;6.25MG;6.25MG** **A217027 005** Jan 23, 2023

AB1 **7.5MG;7.5MG;7.5MG;7.5MG** **A217027 006** Jan 23, 2023

AB1 IMPAX LABS **1.25MG;1.25MG;1.25MG;1.25MG** **A076852 001** Feb 16, 2016

AB1 **2.5MG;2.5MG;2.5MG;2.5MG** **A076852 002** Feb 16, 2016

AB1 **3.75MG;3.75MG;3.75MG;3.75MG** **A076852 003** Feb 16, 2016

AB1 **5MG;5MG;5MG;5MG** **A076852 004** Feb 16, 2016

AB1 **6.25MG;6.25MG;6.25MG;6.25MG** **A076852 005** Feb 16, 2016

AB1 **7.5MG;7.5MG;7.5MG;7.5MG** **A076852 006** Feb 16, 2016

AB1 LANNETT CO INC **1.25MG;1.25MG;1.25MG;1.25MG** **A214403 001** Nov 26, 2021

AB1 **2.5MG;2.5MG;2.5MG;2.5MG** **A214403 002** Nov 26, 2021

PRESCRIPTION DRUG PRODUCT LIST

AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATE

CAPSULE, EXTENDED RELEASE; ORAL

DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE

<u>AB1</u>		<u>3.75MG; 3.75MG; 3.75MG; 3.75MG</u>	<u>A214403 003</u>	Nov 26, 2021
<u>AB1</u>		<u>5MG; 5MG; 5MG; 5MG</u>	<u>A214403 004</u>	Nov 26, 2021
<u>AB1</u>		<u>6.25MG; 6.25MG; 6.25MG; 6.25MG</u>	<u>A214403 005</u>	Nov 26, 2021
<u>AB1</u>		<u>7.5MG; 7.5MG; 7.5MG; 7.5MG</u>	<u>A214403 006</u>	Nov 26, 2021
<u>AB1</u>	LUPIN	<u>1.25MG; 1.25MG; 1.25MG; 1.25MG</u>	<u>A214777 001</u>	Nov 07, 2024
<u>AB1</u>		<u>2.5MG; 2.5MG; 2.5MG; 2.5MG</u>	<u>A214777 002</u>	Nov 07, 2024
<u>AB1</u>		<u>3.75MG; 3.75MG; 3.75MG; 3.75MG</u>	<u>A214777 003</u>	Nov 07, 2024
<u>AB1</u>		<u>5MG; 5MG; 5MG; 5MG</u>	<u>A214777 004</u>	Nov 07, 2024
<u>AB1</u>		<u>6.25MG; 6.25MG; 6.25MG; 6.25MG</u>	<u>A214777 005</u>	Nov 07, 2024
<u>AB1</u>		<u>7.5MG; 7.5MG; 7.5MG; 7.5MG</u>	<u>A214777 006</u>	Nov 07, 2024
<u>AB1</u>	RHODES PHARMS	<u>1.25MG; 1.25MG; 1.25MG; 1.25MG</u>	<u>A210651 001</u>	May 17, 2019
<u>AB1</u>		<u>2.5MG; 2.5MG; 2.5MG; 2.5MG</u>	<u>A210651 002</u>	May 17, 2019
<u>AB1</u>		<u>3.75MG; 3.75MG; 3.75MG; 3.75MG</u>	<u>A210651 003</u>	May 17, 2019
<u>AB1</u>		<u>5MG; 5MG; 5MG; 5MG</u>	<u>A210651 004</u>	May 17, 2019
<u>AB1</u>		<u>6.25MG; 6.25MG; 6.25MG; 6.25MG</u>	<u>A210651 005</u>	May 17, 2019
<u>AB1</u>		<u>7.5MG; 7.5MG; 7.5MG; 7.5MG</u>	<u>A210651 006</u>	May 17, 2019
<u>AB1</u>	SPECGX LLC	<u>1.25MG; 1.25MG; 1.25MG; 1.25MG</u>	<u>A211547 001</u>	Apr 22, 2019
<u>AB1</u>		<u>2.5MG; 2.5MG; 2.5MG; 2.5MG</u>	<u>A211547 002</u>	Apr 22, 2019
<u>AB1</u>		<u>3.75MG; 3.75MG; 3.75MG; 3.75MG</u>	<u>A211547 003</u>	Apr 22, 2019
<u>AB1</u>		<u>5MG; 5MG; 5MG; 5MG</u>	<u>A211547 004</u>	Apr 22, 2019
<u>AB1</u>		<u>6.25MG; 6.25MG; 6.25MG; 6.25MG</u>	<u>A211547 005</u>	Apr 22, 2019
<u>AB1</u>		<u>7.5MG; 7.5MG; 7.5MG; 7.5MG</u>	<u>A211547 006</u>	Apr 22, 2019
<u>AB2</u>		<u>3.125MG; 3.125MG; 3.125MG; 3.125MG</u>	<u>A211546 001</u>	Aug 31, 2023
<u>AB2</u>		<u>6.25MG; 6.25MG; 6.25MG; 6.25MG</u>	<u>A211546 002</u>	Aug 31, 2023
<u>AB2</u>		<u>9.375MG; 9.375MG; 9.375MG; 9.375MG</u>	<u>A211546 003</u>	Aug 31, 2023
<u>AB2</u>		<u>12.5MG; 12.5MG; 12.5MG; 12.5MG</u>	<u>A211546 004</u>	Aug 31, 2023
<u>AB2</u>	SUN PHARM INDS INC	<u>3.125MG; 3.125MG; 3.125MG; 3.125MG</u>	<u>A215997 001</u>	Sep 27, 2023
<u>AB2</u>		<u>6.25MG; 6.25MG; 6.25MG; 6.25MG</u>	<u>A215997 002</u>	Sep 27, 2023
<u>AB2</u>		<u>9.375MG; 9.375MG; 9.375MG; 9.375MG</u>	<u>A215997 003</u>	Sep 27, 2023
<u>AB2</u>		<u>12.5MG; 12.5MG; 12.5MG; 12.5MG</u>	<u>A215997 004</u>	Sep 27, 2023
<u>AB2</u>	TEVA PHARMS USA	<u>3.125MG; 3.125MG; 3.125MG; 3.125MG</u>	<u>A210876 001</u>	Jan 31, 2022
<u>AB2</u>		<u>6.25MG; 6.25MG; 6.25MG; 6.25MG</u>	<u>A210876 002</u>	Jan 31, 2022
<u>AB2</u>		<u>9.375MG; 9.375MG; 9.375MG; 9.375MG</u>	<u>A210876 003</u>	Jan 31, 2022
<u>AB2</u>		<u>12.5MG; 12.5MG; 12.5MG; 12.5MG</u>	<u>A210876 004</u>	Jan 31, 2022
<u>MYDAYIS</u>				
<u>AB2</u>	+	<u>TAKEDA PHARMS USA</u>	<u>3.125MG; 3.125MG; 3.125MG; 3.125MG</u>	<u>N022063 001</u> Jun 20, 2017
<u>AB2</u>	+		<u>6.25MG; 6.25MG; 6.25MG; 6.25MG</u>	<u>N022063 002</u> Jun 20, 2017
<u>AB2</u>	+		<u>9.375MG; 9.375MG; 9.375MG; 9.375MG</u>	<u>N022063 003</u> Jun 20, 2017
<u>AB2</u>	+		<u>12.5MG; 12.5MG; 12.5MG; 12.5MG</u>	<u>N022063 004</u> Jun 20, 2017

TABLET; ORAL

DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE

<u>AB</u>	ACTAVIS ELIZABETH	<u>1.25MG; 1.25MG; 1.25MG; 1.25MG</u>	<u>A206340 001</u>	Feb 05, 2016
<u>AB</u>		<u>1.875MG; 1.875MG; 1.875MG; 1.875MG</u>	<u>A206340 002</u>	Feb 05, 2016
<u>AB</u>		<u>2.5MG; 2.5MG; 2.5MG; 2.5MG</u>	<u>A206340 003</u>	Feb 05, 2016
<u>AB</u>		<u>3.125MG; 3.125MG; 3.125MG; 3.125MG</u>	<u>A206340 004</u>	Feb 05, 2016
<u>AB</u>		<u>3.75MG; 3.75MG; 3.75MG; 3.75MG</u>	<u>A206340 005</u>	Feb 05, 2016
<u>AB</u>		<u>5MG; 5MG; 5MG; 5MG</u>	<u>A206340 006</u>	Feb 05, 2016
<u>AB</u>		<u>7.5MG; 7.5MG; 7.5MG; 7.5MG</u>	<u>A206340 007</u>	Feb 05, 2016
<u>AB</u>	ALKEM LABS LTD	<u>1.25MG; 1.25MG; 1.25MG; 1.25MG</u>	<u>A217787 001</u>	Oct 06, 2023
<u>AB</u>		<u>1.875MG; 1.875MG; 1.875MG; 1.875MG</u>	<u>A217787 002</u>	Oct 06, 2023
<u>AB</u>		<u>2.5MG; 2.5MG; 2.5MG; 2.5MG</u>	<u>A217787 003</u>	Oct 06, 2023
<u>AB</u>		<u>3.125MG; 3.125MG; 3.125MG; 3.125MG</u>	<u>A217787 004</u>	Oct 06, 2023
<u>AB</u>		<u>3.75MG; 3.75MG; 3.75MG; 3.75MG</u>	<u>A217787 005</u>	Oct 06, 2023
<u>AB</u>		<u>5MG; 5MG; 5MG; 5MG</u>	<u>A217787 006</u>	Oct 06, 2023
<u>AB</u>		<u>7.5MG; 7.5MG; 7.5MG; 7.5MG</u>	<u>A217787 007</u>	Oct 06, 2023
<u>AB</u>	ALVOGEN	<u>1.25MG; 1.25MG; 1.25MG; 1.25MG</u>	<u>A207388 001</u>	Jul 28, 2017
<u>AB</u>		<u>1.875MG; 1.875MG; 1.875MG; 1.875MG</u>	<u>A207388 002</u>	Jul 28, 2017
<u>AB</u>		<u>2.5MG; 2.5MG; 2.5MG; 2.5MG</u>	<u>A207388 003</u>	Jul 28, 2017
<u>AB</u>		<u>3.125MG; 3.125MG; 3.125MG; 3.125MG</u>	<u>A207388 004</u>	Jul 28, 2017
<u>AB</u>		<u>3.75MG; 3.75MG; 3.75MG; 3.75MG</u>	<u>A207388 005</u>	Jul 28, 2017
<u>AB</u>		<u>5MG; 5MG; 5MG; 5MG</u>	<u>A207388 006</u>	Jul 28, 2017
<u>AB</u>		<u>7.5MG; 7.5MG; 7.5MG; 7.5MG</u>	<u>A207388 007</u>	Jul 28, 2017
<u>AB</u>	ASCENT PHARMS INC	<u>1.25MG; 1.25MG; 1.25MG; 1.25MG</u>	<u>A213709 001</u>	Apr 22, 2021
<u>AB</u>		<u>1.875MG; 1.875MG; 1.875MG; 1.875MG</u>	<u>A213709 002</u>	Apr 22, 2021
<u>AB</u>		<u>2.5MG; 2.5MG; 2.5MG; 2.5MG</u>	<u>A213709 003</u>	Apr 22, 2021
<u>AB</u>		<u>3.125MG; 3.125MG; 3.125MG; 3.125MG</u>	<u>A213709 004</u>	Apr 22, 2021
<u>AB</u>		<u>3.75MG; 3.75MG; 3.75MG; 3.75MG</u>	<u>A213709 005</u>	Apr 22, 2021
<u>AB</u>		<u>5MG; 5MG; 5MG; 5MG</u>	<u>A213709 006</u>	Apr 22, 2021
<u>AB</u>		<u>7.5MG; 7.5MG; 7.5MG; 7.5MG</u>	<u>A213709 007</u>	Apr 22, 2021

PRESCRIPTION DRUG PRODUCT LIST

AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATE

TABLET;ORAL

DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE

AB	AUROLIFE PHARMA LLC	<u>1.25MG;1.25MG;1.25MG;1.25MG</u>	<u>A202424 001</u>	Nov 27, 2013
AB		<u>1.875MG;1.875MG;1.875MG;1.875MG</u>	<u>A202424 002</u>	Nov 27, 2013
AB		<u>2.5MG;2.5MG;2.5MG;2.5MG</u>	<u>A202424 003</u>	Nov 27, 2013
AB		<u>3.125MG;3.125MG;3.125MG;3.125MG</u>	<u>A202424 004</u>	Nov 27, 2013
AB		<u>3.75MG;3.75MG;3.75MG;3.75MG</u>	<u>A202424 005</u>	Nov 27, 2013
AB		<u>5MG;5MG;5MG;5MG</u>	<u>A202424 006</u>	Nov 27, 2013
AB		<u>7.5MG;7.5MG;7.5MG;7.5MG</u>	<u>A202424 007</u>	Nov 27, 2013
AB	BARR	<u>1.25MG;1.25MG;1.25MG;1.25MG</u>	<u>A040422 001</u>	Feb 11, 2002
AB		<u>1.875MG;1.875MG;1.875MG;1.875MG</u>	<u>A040422 005</u>	Mar 19, 2003
AB		<u>2.5MG;2.5MG;2.5MG;2.5MG</u>	<u>A040422 002</u>	Feb 11, 2002
AB		<u>3.125MG;3.125MG;3.125MG;3.125MG</u>	<u>A040422 006</u>	Mar 19, 2003
AB		<u>3.75MG;3.75MG;3.75MG;3.75MG</u>	<u>A040422 007</u>	Mar 19, 2003
AB		<u>5MG;5MG;5MG;5MG</u>	<u>A040422 003</u>	Feb 11, 2002
AB		<u>7.5MG;7.5MG;7.5MG;7.5MG</u>	<u>A040422 004</u>	Feb 11, 2002
AB	ELITE LABS INC	<u>1.25MG;1.25MG;1.25MG;1.25MG</u>	<u>A211352 001</u>	Dec 07, 2018
AB		<u>1.875MG;1.875MG;1.875MG;1.875MG</u>	<u>A211352 002</u>	Dec 07, 2018
AB		<u>2.5MG;2.5MG;2.5MG;2.5MG</u>	<u>A211352 003</u>	Dec 07, 2018
AB		<u>3.125MG;3.125MG;3.125MG;3.125MG</u>	<u>A211352 004</u>	Dec 07, 2018
AB		<u>3.75MG;3.75MG;3.75MG;3.75MG</u>	<u>A211352 005</u>	Dec 07, 2018
AB		<u>5MG;5MG;5MG;5MG</u>	<u>A211352 006</u>	Dec 07, 2018
AB		<u>7.5MG;7.5MG;7.5MG;7.5MG</u>	<u>A211352 007</u>	Dec 07, 2018
AB	EPIC PHARMA LLC	<u>1.25MG;1.25MG;1.25MG;1.25MG</u>	<u>A040444 001</u>	Jun 19, 2002
AB		<u>1.875MG;1.875MG;1.875MG;1.875MG</u>	<u>A040444 005</u>	Nov 03, 2014
AB		<u>2.5MG;2.5MG;2.5MG;2.5MG</u>	<u>A040444 002</u>	Jun 19, 2002
AB		<u>3.125MG;3.125MG;3.125MG;3.125MG</u>	<u>A040444 006</u>	Nov 03, 2014
AB		<u>3.75MG;3.75MG;3.75MG;3.75MG</u>	<u>A040444 007</u>	Nov 03, 2014
AB		<u>5MG;5MG;5MG;5MG</u>	<u>A040444 003</u>	Jun 19, 2002
AB		<u>7.5MG;7.5MG;7.5MG;7.5MG</u>	<u>A040444 004</u>	Jun 19, 2002
AB	GRANULES	<u>1.25MG;1.25MG;1.25MG;1.25MG</u>	<u>A215771 001</u>	Dec 28, 2021
AB		<u>1.875MG;1.875MG;1.875MG;1.875MG</u>	<u>A215771 002</u>	Dec 28, 2021
AB		<u>2.5MG;2.5MG;2.5MG;2.5MG</u>	<u>A215771 003</u>	Dec 28, 2021
AB		<u>3.125MG;3.125MG;3.125MG;3.125MG</u>	<u>A215771 004</u>	Dec 28, 2021
AB		<u>3.75MG;3.75MG;3.75MG;3.75MG</u>	<u>A215771 005</u>	Dec 28, 2021
AB		<u>5MG;5MG;5MG;5MG</u>	<u>A215771 006</u>	Dec 28, 2021
AB		<u>7.5MG;7.5MG;7.5MG;7.5MG</u>	<u>A215771 007</u>	Dec 28, 2021
AB	LANNETT CO INC	<u>1.25MG;1.25MG;1.25MG;1.25MG</u>	<u>A215565 001</u>	Jul 08, 2022
AB		<u>1.875MG;1.875MG;1.875MG;1.875MG</u>	<u>A215565 002</u>	Jul 08, 2022
AB		<u>2.5MG;2.5MG;2.5MG;2.5MG</u>	<u>A215565 003</u>	Jul 08, 2022
AB		<u>3.125MG;3.125MG;3.125MG;3.125MG</u>	<u>A215565 004</u>	Jul 08, 2022
AB		<u>3.75MG;3.75MG;3.75MG;3.75MG</u>	<u>A215565 005</u>	Jul 08, 2022
AB		<u>5MG;5MG;5MG;5MG</u>	<u>A215565 006</u>	Jul 08, 2022
AB		<u>7.5MG;7.5MG;7.5MG;7.5MG</u>	<u>A215565 007</u>	Jul 08, 2022
AB	NUVO PHARM	<u>1.25MG;1.25MG;1.25MG;1.25MG</u>	<u>A209799 001</u>	Dec 28, 2017
AB		<u>1.875MG;1.875MG;1.875MG;1.875MG</u>	<u>A209799 002</u>	Dec 28, 2017
AB		<u>2.5MG;2.5MG;2.5MG;2.5MG</u>	<u>A209799 003</u>	Dec 28, 2017
AB		<u>3.125MG;3.125MG;3.125MG;3.125MG</u>	<u>A209799 004</u>	Dec 28, 2017
AB		<u>3.75MG;3.75MG;3.75MG;3.75MG</u>	<u>A209799 005</u>	Dec 28, 2017
AB		<u>5MG;5MG;5MG;5MG</u>	<u>A209799 006</u>	Dec 28, 2017
AB		<u>7.5MG;7.5MG;7.5MG;7.5MG</u>	<u>A209799 007</u>	Dec 28, 2017
AB	ORYZA	<u>1.25MG;1.25MG;1.25MG;1.25MG</u>	<u>A210293 001</u>	Apr 03, 2020
AB		<u>1.875MG;1.875MG;1.875MG;1.875MG</u>	<u>A210293 002</u>	Apr 03, 2020
AB		<u>2.5MG;2.5MG;2.5MG;2.5MG</u>	<u>A210293 003</u>	Apr 03, 2020
AB		<u>3.75MG;3.75MG;3.75MG;3.75MG</u>	<u>A210293 004</u>	Apr 03, 2020
AB		<u>5MG;5MG;5MG;5MG</u>	<u>A210293 005</u>	Apr 03, 2020
AB		<u>7.5MG;7.5MG;7.5MG;7.5MG</u>	<u>A210293 006</u>	Apr 03, 2020
AB	RHODES PHARMS	<u>1.25MG;1.25MG;1.25MG;1.25MG</u>	<u>A213111 001</u>	Jan 13, 2021
AB		<u>1.875MG;1.875MG;1.875MG;1.875MG</u>	<u>A213111 002</u>	Jan 13, 2021
AB		<u>2.5MG;2.5MG;2.5MG;2.5MG</u>	<u>A213111 003</u>	Jan 13, 2021
AB		<u>3.125MG;3.125MG;3.125MG;3.125MG</u>	<u>A213111 004</u>	Jan 13, 2021
AB		<u>3.75MG;3.75MG;3.75MG;3.75MG</u>	<u>A213111 005</u>	Jan 13, 2021
AB		<u>5MG;5MG;5MG;5MG</u>	<u>A213111 006</u>	Jan 13, 2021
AB		<u>7.5MG;7.5MG;7.5MG;7.5MG</u>	<u>A213111 007</u>	Jan 13, 2021
AB	SANDOZ	<u>1.25MG;1.25MG;1.25MG;1.25MG</u>	<u>A040439 004</u>	Sep 27, 2002
AB		<u>2.5MG;2.5MG;2.5MG;2.5MG</u>	<u>A040439 001</u>	Jun 14, 2002
AB		<u>5MG;5MG;5MG;5MG</u>	<u>A040439 002</u>	Jun 14, 2002
AB		<u>7.5MG;7.5MG;7.5MG;7.5MG</u>	<u>A040439 003</u>	Jun 14, 2002
AB	SPECGX LLC	<u>1.25MG;1.25MG;1.25MG;1.25MG</u>	<u>A040440 001</u>	Oct 07, 2003
AB		<u>1.875MG;1.875MG;1.875MG;1.875MG</u>	<u>A040440 002</u>	Oct 07, 2003
AB		<u>2.5MG;2.5MG;2.5MG;2.5MG</u>	<u>A040440 003</u>	Oct 07, 2003

PRESCRIPTION DRUG PRODUCT LIST

AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATE

TABLET; ORAL

DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE

AB		3.125MG;3.125MG;3.125MG;3.125MG	A040440 004	Oct 07, 2003
AB		3.75MG;3.75MG;3.75MG;3.75MG	A040440 005	Oct 07, 2003
AB		5MG;5MG;5MG;5MG	A040440 006	Oct 07, 2003
AB		7.5MG;7.5MG;7.5MG;7.5MG	A040440 007	Oct 07, 2003
AB	SUN PHARM INDUSTRIES	1.25MG;1.25MG;1.25MG;1.25MG	A040480 001	Sep 09, 2003
AB		1.875MG;1.875MG;1.875MG;1.875MG	A040480 002	Sep 09, 2003
AB		2.5MG;2.5MG;2.5MG;2.5MG	A040480 003	Sep 09, 2003
AB		3.125MG;3.125MG;3.125MG;3.125MG	A040480 004	Sep 09, 2003
AB		3.75MG;3.75MG;3.75MG;3.75MG	A040480 005	Sep 09, 2003
AB		5MG;5MG;5MG;5MG	A040480 006	Sep 09, 2003
AB		7.5MG;7.5MG;7.5MG;7.5MG	A040480 007	Sep 09, 2003
AB	ZYDUS PHARMS	1.25MG;1.25MG;1.25MG;1.25MG	A207340 001	Oct 31, 2017
AB		1.875MG;1.875MG;1.875MG;1.875MG	A207340 002	Oct 31, 2017
AB		2.5MG;2.5MG;2.5MG;2.5MG	A207340 003	Oct 31, 2017
AB		3.125MG;3.125MG;3.125MG;3.125MG	A207340 004	Oct 31, 2017
AB		3.75MG;3.75MG;3.75MG;3.75MG	A207340 005	Oct 31, 2017
AB		5MG;5MG;5MG;5MG	A207340 006	Oct 31, 2017
AB		7.5MG;7.5MG;7.5MG;7.5MG	A207340 007	Oct 31, 2017

AMPHETAMINE SULFATE

TABLET; ORAL

AMPHETAMINE SULFATE

AA	ALKEM LABS LTD	5MG	A213720 001	Oct 27, 2020
AA		10MG	A213720 002	Oct 27, 2020
AA	AMNEAL PHARMS	5MG	A211139 001	Sep 26, 2018
AA		10MG	A211139 002	Sep 26, 2018
AA	AUROLIFE PHARMA LLC	5MG	A211639 001	Apr 17, 2019
AA		10MG	A211639 002	Apr 17, 2019
AA	BIONPHARMA	5MG	A212919 001	Nov 22, 2019
AA		10MG	A212919 002	Nov 22, 2019
AA	EPIC PHARMA LLC	5MG	A213980 001	Oct 27, 2020
AA		10MG	A213980 002	Oct 27, 2020
AA	GRANULES	5MG	A212619 001	Aug 05, 2019
AA		10MG	A212619 002	Aug 05, 2019
AA	PRINSTON INC	5MG	A211861 001	Mar 11, 2020
AA		10MG	A211861 002	Mar 11, 2020
AA	RHODES PHARMS	5MG	A213852 001	Sep 07, 2021
AA		10MG	A213852 002	Sep 07, 2021
AA	SANALUZ	5MG	A212582 001	Feb 04, 2020
AA		10MG	A212582 002	Feb 04, 2020
AA	SENORES PHARMS	5MG	A212901 001	May 22, 2020
AA		10MG	A212901 002	May 22, 2020
AA	SPECGX LLC	5MG	A213583 001	Jan 22, 2021
AA		10MG	A213583 002	Jan 22, 2021
	EVEKEO			
AA	AZURITY	5MG	A200166 001	Aug 09, 2012
AA	!	10MG	A200166 002	Aug 09, 2012

AMPHETAMINE; AMPHETAMINE ASPARTATE/DEXTROAMPHETAMINE SULFATE

SUSPENSION, EXTENDED RELEASE; ORAL

DYANAVEL XR

+! TRIS PHARMA INC 2MG/ML;EQ 0.5MG BASE/ML N208147 001 Oct 19, 2015

TABLET, EXTENDED RELEASE; ORAL

DYANAVEL XR 10

+ TRIS PHARMA INC 8MG;EQ 2MG BASE N210526 002 Nov 04, 2021

DYANAVEL XR 15

+ TRIS PHARMA INC 12MG;EQ 3MG BASE N210526 003 Nov 04, 2021

DYANAVEL XR 20

+! TRIS PHARMA INC 16MG;EQ 4MG BASE N210526 004 Nov 04, 2021

DYANAVEL XR 5

+ TRIS PHARMA INC 4MG;EQ 1MG BASE N210526 001 Nov 04, 2021

AMPHOTERICIN B

INJECTABLE; INJECTION

AMPHOTERICIN B

! XGEN PHARMS 50MG/VIAL A063206 001 Apr 29, 1992

INJECTABLE, LIPID COMPLEX; INJECTION

ABELCET

+! LEADIANT BIOSCI INC 5MG/ML N050724 001 Nov 20, 1995

PRESCRIPTION DRUG PRODUCT LIST

AMPHOTERICIN B

INJECTABLE, LIPOSOMAL; INJECTION

AMBISOME

<u>AB</u>	<u>+</u> !	ASTELLAS	<u>50MG/VIAL</u>	<u>N050740</u>	<u>001</u>	Aug 11, 1997
-----------	------------	----------	------------------	----------------	------------	--------------

AMPHOTERICIN B

<u>AB</u>		EUGIA PHARMA	<u>50MG/VIAL</u>	<u>A214010</u>	<u>001</u>	Nov 17, 2022
-----------	--	--------------	------------------	----------------	------------	--------------

<u>AB</u>		SPIL	<u>50MG/VIAL</u>	<u>A212514</u>	<u>001</u>	Dec 14, 2021
-----------	--	------	------------------	----------------	------------	--------------

AMPICILLIN SODIUM

INJECTABLE; INJECTION

AMPICILLIN SODIUM

<u>AP</u>		ACS DOBFAR SPA	<u>EQ 10GM BASE/VIAL</u>	<u>A090889</u>	<u>001</u>	Apr 03, 2013
-----------	--	----------------	--------------------------	----------------	------------	--------------

<u>AP</u>		ANTIBIOTICE	<u>EQ 250MG BASE/VIAL</u>	<u>A090354</u>	<u>001</u>	Dec 28, 2009
-----------	--	-------------	---------------------------	----------------	------------	--------------

<u>AP</u>			<u>EQ 500MG BASE/VIAL</u>	<u>A090354</u>	<u>002</u>	Dec 28, 2009
-----------	--	--	---------------------------	----------------	------------	--------------

<u>AP</u>			<u>EQ 1GM BASE/VIAL</u>	<u>A090354</u>	<u>003</u>	Dec 28, 2009
-----------	--	--	-------------------------	----------------	------------	--------------

<u>AP</u>			<u>EQ 2GM BASE/VIAL</u>	<u>A090354</u>	<u>004</u>	Dec 28, 2009
-----------	--	--	-------------------------	----------------	------------	--------------

<u>AP</u>		EUGIA PHARMA	<u>EQ 250MG BASE/VIAL</u>	<u>A065499</u>	<u>002</u>	Aug 17, 2010
-----------	--	--------------	---------------------------	----------------	------------	--------------

<u>AP</u>		SPECLTS				
-----------	--	---------	--	--	--	--

<u>AP</u>			<u>EQ 500MG BASE/VIAL</u>	<u>A065499</u>	<u>003</u>	Aug 17, 2010
-----------	--	--	---------------------------	----------------	------------	--------------

<u>AP</u>			<u>EQ 1GM BASE/VIAL</u>	<u>A065499</u>	<u>004</u>	Aug 17, 2010
-----------	--	--	-------------------------	----------------	------------	--------------

<u>AP</u>			<u>EQ 2GM BASE/VIAL</u>	<u>A065499</u>	<u>005</u>	Aug 17, 2010
-----------	--	--	-------------------------	----------------	------------	--------------

<u>AP</u>			<u>EQ 10GM BASE/VIAL</u>	<u>A065493</u>	<u>001</u>	Aug 17, 2010
-----------	--	--	--------------------------	----------------	------------	--------------

<u>AP</u>		HQ SPECLT PHARMA	<u>EQ 250MG BASE/VIAL</u>	<u>A062772</u>	<u>006</u>	Apr 15, 1993
-----------	--	------------------	---------------------------	----------------	------------	--------------

<u>AP</u>			<u>EQ 500MG BASE/VIAL</u>	<u>A062772</u>	<u>007</u>	Apr 15, 1993
-----------	--	--	---------------------------	----------------	------------	--------------

<u>AP</u>			<u>EQ 1GM BASE/VIAL</u>	<u>A062772</u>	<u>001</u>	Apr 15, 1993
-----------	--	--	-------------------------	----------------	------------	--------------

<u>AP</u>			<u>EQ 2GM BASE/VIAL</u>	<u>A062772</u>	<u>003</u>	Apr 15, 1993
-----------	--	--	-------------------------	----------------	------------	--------------

<u>AP</u>			<u>EQ 10GM BASE/VIAL</u>	<u>A063142</u>	<u>001</u>	Apr 15, 1993
-----------	--	--	--------------------------	----------------	------------	--------------

<u>AP</u>		ISTITUTO BIO ITA	<u>EQ 10GM BASE/VIAL</u>	<u>A201404</u>	<u>001</u>	Dec 20, 2013
-----------	--	------------------	--------------------------	----------------	------------	--------------

<u>AP</u>		SPA				
-----------	--	-----	--	--	--	--

<u>AP</u>			<u>EQ 250MG BASE/VIAL</u>	<u>A062719</u>	<u>001</u>	May 12, 1987
-----------	--	--	---------------------------	----------------	------------	--------------

<u>AP</u>			<u>EQ 500MG BASE/VIAL</u>	<u>A062719</u>	<u>003</u>	May 12, 1987
-----------	--	--	---------------------------	----------------	------------	--------------

<u>AP</u>			<u>EQ 1GM BASE/VIAL</u>	<u>A062719</u>	<u>002</u>	May 12, 1987
-----------	--	--	-------------------------	----------------	------------	--------------

<u>AP</u>			<u>EQ 2GM BASE/VIAL</u>	<u>A062797</u>	<u>002</u>	Jul 12, 1993
-----------	--	--	-------------------------	----------------	------------	--------------

<u>AP</u>		SAGENT PHARMS INC	<u>EQ 125MG BASE/VIAL</u>	<u>A090583</u>	<u>001</u>	Nov 27, 2015
-----------	--	-------------------	---------------------------	----------------	------------	--------------

<u>AP</u>			<u>EQ 250MG BASE/VIAL</u>	<u>A090583</u>	<u>002</u>	Nov 27, 2015
-----------	--	--	---------------------------	----------------	------------	--------------

<u>AP</u>			<u>EQ 500MG BASE/VIAL</u>	<u>A090583</u>	<u>003</u>	Nov 27, 2015
-----------	--	--	---------------------------	----------------	------------	--------------

<u>AP</u>			<u>EQ 1GM BASE/VIAL</u>	<u>A090583</u>	<u>004</u>	Nov 27, 2015
-----------	--	--	-------------------------	----------------	------------	--------------

<u>AP</u>			<u>EQ 2GM BASE/VIAL</u>	<u>A090583</u>	<u>005</u>	Nov 27, 2015
-----------	--	--	-------------------------	----------------	------------	--------------

<u>AP</u>			<u>EQ 10GM BASE/VIAL</u>	<u>A090581</u>	<u>001</u>	Oct 20, 2015
-----------	--	--	--------------------------	----------------	------------	--------------

<u>AP</u>	<u>!</u>	SANDOZ	<u>EQ 125MG BASE/VIAL</u>	<u>A061395</u>	<u>001</u>	
-----------	----------	--------	---------------------------	----------------	------------	--

<u>AP</u>	<u>!</u>		<u>EQ 250MG BASE/VIAL</u>	<u>A061395</u>	<u>002</u>	
-----------	----------	--	---------------------------	----------------	------------	--

<u>AP</u>	<u>!</u>		<u>EQ 500MG BASE/VIAL</u>	<u>A061395</u>	<u>003</u>	
-----------	----------	--	---------------------------	----------------	------------	--

<u>AP</u>	<u>!</u>		<u>EQ 1GM BASE/VIAL</u>	<u>A061395</u>	<u>004</u>	
-----------	----------	--	-------------------------	----------------	------------	--

<u>AP</u>	<u>!</u>		<u>EQ 2GM BASE/VIAL</u>	<u>A061395</u>	<u>005</u>	
-----------	----------	--	-------------------------	----------------	------------	--

<u>AP</u>	<u>!</u>		<u>EQ 10GM BASE/VIAL</u>	<u>A061395</u>	<u>006</u>	
-----------	----------	--	--------------------------	----------------	------------	--

<u>AP</u>		STERISCIENCE	<u>EQ 1GM BASE/VIAL</u>	<u>A201025</u>	<u>003</u>	Apr 09, 2014
-----------	--	--------------	-------------------------	----------------	------------	--------------

<u>AP</u>		SPECLTS				
-----------	--	---------	--	--	--	--

<u>AP</u>			<u>EQ 2GM BASE/VIAL</u>	<u>A201025</u>	<u>004</u>	Apr 09, 2014
-----------	--	--	-------------------------	----------------	------------	--------------

<u>AP</u>			<u>EQ 10GM BASE/VIAL</u>	<u>A202198</u>	<u>001</u>	Apr 07, 2014
-----------	--	--	--------------------------	----------------	------------	--------------

POWDER; INTRAVENOUS

AMPICILLIN SODIUM

<u>AP</u>	<u>!</u>	SANDOZ	<u>EQ 1GM BASE/VIAL</u>	<u>A062738</u>	<u>001</u>	Feb 19, 1987
-----------	----------	--------	-------------------------	----------------	------------	--------------

<u>AP</u>	<u>!</u>		<u>EQ 2GM BASE/VIAL</u>	<u>A062738</u>	<u>002</u>	Feb 19, 1987
-----------	----------	--	-------------------------	----------------	------------	--------------

AMPICILLIN SODIUM; SULBACTAM SODIUM

INJECTABLE; INJECTION

AMPICILLIN AND SULBACTAM

<u>AP</u>		ACS DOBFAR	<u>EQ 1GM BASE/VIAL;EQ 500MG BASE/VIAL</u>	<u>A065406</u>	<u>001</u>	Dec 22, 2009
-----------	--	------------	--	----------------	------------	--------------

<u>AP</u>			<u>EQ 2GM BASE/VIAL;EQ 1GM BASE/VIAL</u>	<u>A065406</u>	<u>002</u>	Dec 22, 2009
-----------	--	--	--	----------------	------------	--------------

<u>AP</u>			<u>EQ 10GM BASE/VIAL;EQ 5GM BASE/VIAL</u>	<u>A065403</u>	<u>001</u>	Dec 23, 2009
-----------	--	--	---	----------------	------------	--------------

<u>AP</u>		ANTIBIOTICE	<u>EQ 1GM BASE/VIAL;EQ 500MG BASE/VIAL</u>	<u>A201406</u>	<u>001</u>	Dec 07, 2015
-----------	--	-------------	--	----------------	------------	--------------

<u>AP</u>			<u>EQ 2GM BASE/VIAL;EQ 1GM BASE/VIAL</u>	<u>A201406</u>	<u>002</u>	Dec 07, 2015
-----------	--	--	--	----------------	------------	--------------

<u>AP</u>		EUGIA PHARMA	<u>EQ 1GM BASE/VIAL;EQ 500MG BASE/VIAL</u>	<u>A090349</u>	<u>001</u>	Sep 20, 2010
-----------	--	--------------	--	----------------	------------	--------------

<u>AP</u>		SPECLTS				
-----------	--	---------	--	--	--	--

<u>AP</u>			<u>EQ 2GM BASE/VIAL;EQ 1GM BASE/VIAL</u>	<u>A090349</u>	<u>002</u>	Sep 20, 2010
-----------	--	--	--	----------------	------------	--------------

<u>AP</u>			<u>EQ 10GM BASE/VIAL;EQ 5GM BASE/VIAL</u>	<u>A090339</u>	<u>001</u>	Sep 20, 2010
-----------	--	--	---	----------------	------------	--------------

<u>AP</u>		HIKMA	<u>EQ 1GM BASE/VIAL;EQ 500MG BASE/VIAL</u>	<u>A065074</u>	<u>001</u>	Mar 19, 2002
-----------	--	-------	--	----------------	------------	--------------

<u>AP</u>			<u>EQ 2GM BASE/VIAL;EQ 1GM BASE/VIAL</u>	<u>A065074</u>	<u>002</u>	Mar 19, 2002
-----------	--	--	--	----------------	------------	--------------

<u>AP</u>			<u>EQ 10GM BASE/VIAL;EQ 5GM BASE/VIAL</u>	<u>A065076</u>	<u>001</u>	Mar 19, 2002
-----------	--	--	---	----------------	------------	--------------

<u>AP</u>		HQ SPECLT PHARMA	<u>EQ 1GM BASE/VIAL;EQ 500MG BASE/VIAL</u>	<u>A065176</u>	<u>001</u>	Nov 30, 2005
-----------	--	------------------	--	----------------	------------	--------------

<u>AP</u>			<u>EQ 2GM BASE/VIAL;EQ 1GM BASE/VIAL</u>	<u>A065176</u>	<u>002</u>	Nov 30, 2005
-----------	--	--	--	----------------	------------	--------------

<u>AP</u>			<u>EQ 10GM BASE/VIAL;EQ 5GM BASE/VIAL</u>	<u>A065188</u>	<u>001</u>	Nov 25, 2005
-----------	--	--	---	----------------	------------	--------------

<u>AP</u>		ISTITUTO BIO ITA	<u>EQ 1GM BASE/VIAL;EQ 500MG BASE/VIAL</u>	<u>A065222</u>	<u>001</u>	Nov 29, 2005
-----------	--	------------------	--	----------------	------------	--------------

<u>AP</u>		SPA				
-----------	--	-----	--	--	--	--

PRESCRIPTION DRUG PRODUCT LIST

AMPICILLIN SODIUM; SULBACTAM SODIUM

INJECTABLE; INJECTION

AMPICILLIN AND SULBACTAM

<u>AP</u>		<u>EQ 2GM BASE/VIAL;EQ 1GM BASE/VIAL</u>	<u>A065222 002</u>	Nov 29, 2005
<u>AP</u>		<u>EQ 10GM BASE/VIAL;EQ 5GM BASE/VIAL</u>	<u>A065314 001</u>	Nov 27, 2006
<u>AP</u>	SANDOZ	<u>EQ 1GM BASE/VIAL;EQ 500MG BASE/VIAL</u>	<u>A065241 001</u>	Jul 25, 2006
<u>AP</u>		<u>EQ 1GM BASE/VIAL;EQ 500MG BASE/VIAL</u>	<u>A065310 001</u>	Jul 25, 2006
<u>AP</u>		<u>EQ 2GM BASE/VIAL;EQ 1GM BASE/VIAL</u>	<u>A065241 002</u>	Jul 25, 2006
<u>AP</u>		<u>EQ 2GM BASE/VIAL;EQ 1GM BASE/VIAL</u>	<u>A065310 002</u>	Jul 25, 2006
<u>AP</u>		<u>EQ 10GM BASE/VIAL;EQ 5GM BASE/VIAL</u>	<u>A065240 001</u>	Jul 25, 2006
<u>AP</u>	STERISCIENCE SPECLTS	<u>EQ 1GM BASE/VIAL;EQ 500MG BASE/VIAL</u>	<u>A201024 001</u>	Apr 07, 2014
<u>AP</u>		<u>EQ 2GM BASE/VIAL;EQ 1GM BASE/VIAL</u>	<u>A201024 002</u>	Apr 07, 2014
<u>AP</u>		<u>EQ 10GM BASE/VIAL;EQ 5GM BASE/VIAL</u>	<u>A202197 001</u>	Apr 07, 2014
<u>UNASYN</u>				
<u>AP</u>	+! PFIZER	<u>EQ 1GM BASE/VIAL;EQ 500MG BASE/VIAL</u>	<u>N050608 002</u>	Dec 31, 1986
<u>AP</u>	+!	<u>EQ 2GM BASE/VIAL;EQ 1GM BASE/VIAL</u>	<u>N050608 001</u>	Dec 31, 1986
<u>AP</u>	+!	<u>EQ 10GM BASE/VIAL;EQ 5GM BASE/VIAL</u>	<u>N050608 005</u>	Dec 10, 1993

AMPICILLIN/AMPICILLIN TRIHYDRATE

CAPSULE; ORAL

AMPICILLIN TRIHYDRATE

<u>AB</u>	AUROBINDO PHARMA	<u>EQ 500MG BASE</u>	<u>A216554 002</u>	Oct 31, 2023
<u>AB</u>	! SANDOZ	<u>EQ 500MG BASE</u>	<u>A064082 002</u>	Aug 29, 1995
	AUROBINDO PHARMA	EQ 250MG BASE	A216554 001	Oct 31, 2023

ANAGRELIDE HYDROCHLORIDE

CAPSULE; ORAL

AGRYLIN

<u>AB</u>	TAKEDA PHARMS USA	<u>EQ 0.5MG BASE</u>	<u>N020333 001</u>	Mar 14, 1997
<u>ANAGRELIDE HYDROCHLORIDE</u>				
<u>AB</u>	IMPAX LABS	<u>EQ 0.5MG BASE</u>	<u>A076910 001</u>	Apr 18, 2005
<u>AB</u>		<u>EQ 1MG BASE</u>	<u>A076910 002</u>	Apr 18, 2005
<u>AB</u>	IVAX SUB TEVA PHARMS	<u>EQ 0.5MG BASE</u>	<u>A076468 001</u>	Apr 18, 2005
<u>AB</u>	!	<u>EQ 1MG BASE</u>	<u>A076468 002</u>	Apr 18, 2005
<u>AB</u>	TORRENT	<u>EQ 0.5MG BASE</u>	<u>A209151 001</u>	Jun 30, 2017
<u>AB</u>		<u>EQ 1MG BASE</u>	<u>A209151 002</u>	Jun 30, 2017

ANASTROZOLE

TABLET; ORAL

ANASTROZOLE

<u>AB</u>	ACCORD HLTHCARE	<u>1MG</u>	<u>A090568 001</u>	Jun 28, 2010
<u>AB</u>	BEIJING YILING	<u>1MG</u>	<u>A206037 001</u>	Nov 09, 2018
<u>AB</u>	CIPLA	<u>1MG</u>	<u>A091164 001</u>	Jun 28, 2010
<u>AB</u>	EUGIA PHARMA	<u>1MG</u>	<u>A212434 001</u>	Jul 24, 2020
<u>AB</u>	KENTON	<u>1MG</u>	<u>A078944 001</u>	Jun 28, 2010
<u>AB</u>	NATCO PHARMA LTD	<u>1MG</u>	<u>A079220 001</u>	Jun 28, 2010
<u>AB</u>	TEVA PHARMS	<u>1MG</u>	<u>A078058 001</u>	Jun 28, 2010
<u>AB</u>	ZYDUS PHARMS USA INC	<u>1MG</u>	<u>A078921 001</u>	Jun 28, 2010
<u>ARIMIDEX</u>				
<u>AB</u>	+! ANI PHARMS	<u>1MG</u>	<u>N020541 001</u>	Dec 27, 1995

ANGIOTENSIN II ACETATE

SOLUTION; INTRAVENOUS

GIAPREZA

+!	LA JOLLA PHARMA	EQ 0.5MG BASE/ML (EQ 0.5MG BASE/ML)	N209360 003	Dec 23, 2021
+!		EQ 2.5MG BASE/ML (EQ 2.5MG BASE/ML)	N209360 001	Dec 21, 2017

ANIDULAFUNGIN

POWDER; INTRAVENOUS

ERAXIS

+!	VICURON HOLDINGS	50MG/VIAL	N021632 001	Feb 17, 2006
+!		100MG/VIAL	N021632 002	Nov 14, 2006

APALUTAMIDE

TABLET; ORAL

ERLEADA

+	JANSSEN BIOTECH	60MG	N210951 001	Feb 14, 2018
+!		240MG	N210951 002	Feb 17, 2023

PRESCRIPTION DRUG PRODUCT LIST

APIXABAN

TABLET; ORAL

APIXABAN

<u>AB</u>	ACCORD HLTHCARE	<u>2.5MG</u>	<u>A210180 001</u>	Jul 28, 2020
<u>AB</u>		<u>5MG</u>	<u>A210180 002</u>	Jul 28, 2020
<u>AB</u>	HETERO LABS LTD V	<u>2.5MG</u>	<u>A210066 001</u>	Nov 21, 2023
<u>AB</u>		<u>5MG</u>	<u>A210066 002</u>	Nov 21, 2023
<u>AB</u>	INDOCO	<u>2.5MG</u>	<u>A209898 001</u>	Sep 11, 2020
<u>AB</u>		<u>5MG</u>	<u>A209898 002</u>	Sep 11, 2020
<u>AB</u>	TORRENT	<u>2.5MG</u>	<u>A210156 001</u>	Dec 17, 2024
<u>AB</u>		<u>5MG</u>	<u>A210156 002</u>	Dec 17, 2024

ELIQUIS

<u>AB</u>	+ BRISTOL MYERS SQUIBB	<u>2.5MG</u>	<u>N202155 001</u>	Dec 28, 2012
<u>AB</u>	+!	<u>5MG</u>	<u>N202155 002</u>	Dec 28, 2012

APOMORPHINE HYDROCHLORIDE

INJECTABLE; SUBCUTANEOUS

APOKYN

<u>AP</u>	+! MDD US	<u>30MG/3ML (10MG/ML)</u>	<u>N021264 002</u>	Apr 20, 2004
-----------	-----------	---------------------------	--------------------	--------------

APOMORPHINE HYDROCHLORIDE

<u>AP</u>	TRUPHARMA	<u>30MG/3ML (10MG/ML)</u>	<u>A212025 001</u>	Feb 23, 2022
-----------	-----------	---------------------------	--------------------	--------------

APRACLONIDINE HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

APRACLONIDINE HYDROCHLORIDE

<u>AT</u>	RISING	<u>EQ 0.5% BASE</u>	<u>A077764 001</u>	Mar 12, 2009
<u>AT</u>	+! HARROW EYE	<u>EQ 0.5% BASE</u>	<u>N020258 001</u>	Jul 30, 1993
	+!	<u>EQ 1% BASE</u>	<u>N019779 001</u>	Dec 31, 1987

APREMILAST

TABLET; ORAL

APREMILAST

<u>AB</u>	ALKEM LABS LTD	<u>10MG</u>	<u>A211761 001</u>	Sep 21, 2021
<u>AB</u>		<u>20MG</u>	<u>A211761 002</u>	Sep 21, 2021
<u>AB</u>		<u>30MG</u>	<u>A211761 003</u>	Sep 21, 2021
<u>AB</u>	ANNORA	<u>10MG</u>	<u>A211878 001</u>	Jul 26, 2023
<u>AB</u>		<u>20MG</u>	<u>A211878 002</u>	Jul 26, 2023
<u>AB</u>		<u>30MG</u>	<u>A211878 003</u>	Jul 26, 2023
<u>AB</u>	GLENMARK PHARMS LTD	<u>10MG</u>	<u>A211674 001</u>	Oct 16, 2023
<u>AB</u>		<u>20MG</u>	<u>A211674 002</u>	Oct 16, 2023
<u>AB</u>		<u>30MG</u>	<u>A211674 003</u>	Oct 16, 2023
<u>AB</u>	MACLEODS PHARMS LTD	<u>10MG</u>	<u>A211788 001</u>	Nov 25, 2024
<u>AB</u>		<u>20MG</u>	<u>A211788 002</u>	Nov 25, 2024
<u>AB</u>		<u>30MG</u>	<u>A211788 003</u>	Nov 25, 2024
<u>AB</u>	MANKIND PHARMA	<u>10MG</u>	<u>A211734 001</u>	Feb 07, 2024
<u>AB</u>		<u>20MG</u>	<u>A211734 002</u>	Feb 07, 2024
<u>AB</u>		<u>30MG</u>	<u>A211734 003</u>	Feb 07, 2024
<u>AB</u>	MSN	<u>10MG</u>	<u>A211887 001</u>	Dec 16, 2024
<u>AB</u>		<u>20MG</u>	<u>A211887 002</u>	Dec 16, 2024
<u>AB</u>		<u>30MG</u>	<u>A211887 003</u>	Dec 16, 2024
<u>AB</u>	SHILPA	<u>10MG</u>	<u>A211774 001</u>	Apr 07, 2023
<u>AB</u>		<u>20MG</u>	<u>A211774 002</u>	Apr 07, 2023
<u>AB</u>		<u>30MG</u>	<u>A211774 003</u>	Apr 07, 2023

OTEZLA

<u>AB</u>	+ AMGEN INC	<u>10MG</u>	<u>N205437 001</u>	Mar 21, 2014
<u>AB</u>	+	<u>20MG</u>	<u>N205437 002</u>	Mar 21, 2014
<u>AB</u>	+!	<u>30MG</u>	<u>N205437 003</u>	Mar 21, 2014

APREPITANT

CAPSULE; ORAL

APREPITANT

<u>AB</u>	GLENMARK SPECLT	<u>40MG</u>	<u>A207777 001</u>	Oct 12, 2017
<u>AB</u>		<u>80MG</u>	<u>A207777 002</u>	Oct 12, 2017
<u>AB</u>		<u>125MG</u>	<u>A207777 003</u>	Oct 12, 2017
<u>AB</u>	SANDOZ	<u>40MG</u>	<u>A090999 001</u>	Sep 24, 2012
<u>AB</u>		<u>80MG</u>	<u>A090999 002</u>	Sep 24, 2012
<u>AB</u>		<u>125MG</u>	<u>A090999 003</u>	Sep 24, 2012
<u>AB</u>	TORRENT	<u>40MG</u>	<u>A211835 001</u>	Oct 21, 2020
<u>AB</u>		<u>80MG</u>	<u>A211835 002</u>	Oct 21, 2020
<u>AB</u>		<u>125MG</u>	<u>A211835 003</u>	Oct 21, 2020

EMEND

<u>AB</u>	+ MERCK	<u>80MG</u>	<u>N021549 001</u>	Mar 26, 2003
<u>AB</u>	+!	<u>125MG</u>	<u>N021549 002</u>	Mar 26, 2003

PRESCRIPTION DRUG PRODUCT LISTAPREPITANT

EMULSION; INTRAVENOUS

APONVIE

+! HERON THERAPS INC 32MG/4.4ML (7.2MG/ML) N216457 001 Sep 16, 2022

CINVANTI

+! HERON THERAPS INC 130MG/18ML (7.2MG/ML) N209296 001 Nov 09, 2017

FOR SUSPENSION; ORAL

EMEND

+! MSD MERCK CO 125MG/KIT N207865 001 Dec 17, 2015

APROCITENTAN

TABLET; ORAL

TRYVIO

+! IDORSIA 12.5MG N217686 001 Mar 19, 2024

ARFORMOTEROL TARTRATE

SOLUTION; INHALATION

ARFORMOTEROL TARTRATE

AN	AIPING PHARM INC	<u>EQ 0.015MG BASE/2ML</u>	<u>A214901 001</u>	Apr 02, 2024
AN	ALEMBIC	<u>EQ 0.015MG BASE/2ML</u>	<u>A214779 001</u>	May 10, 2022
AN	BE PHARMS	<u>EQ 0.015MG BASE/2ML</u>	<u>A213762 001</u>	Jun 22, 2021
AN	CF PHARMTECH	<u>EQ 0.015MG BASE/2ML</u>	<u>A216303 001</u>	May 17, 2024
AN	CIPLA	<u>EQ 0.015MG BASE/2ML</u>	<u>A207306 001</u>	Jun 22, 2021
AN	DR REDDYS	<u>EQ 0.015MG BASE/2ML</u>	<u>A215032 001</u>	Jun 03, 2024
AN	LEXENPHARM	<u>EQ 0.015MG BASE/2ML</u>	<u>A218156 001</u>	Nov 18, 2024
AN	LUPIN	<u>EQ 0.015MG BASE/2ML</u>	<u>A213068 001</u>	Feb 07, 2022
AN	MANKIND PHARMA	<u>EQ 0.015MG BASE/2ML</u>	<u>A216128 001</u>	Nov 15, 2022
AN	RITEDOSE CORP	<u>EQ 0.015MG BASE/2ML</u>	<u>A214736 001</u>	Mar 02, 2022
AN	SUN PHARM	<u>EQ 0.015MG BASE/2ML</u>	<u>A215385 001</u>	May 26, 2022
AN	TEVA PHARMS USA	<u>EQ 0.015MG BASE/2ML</u>	<u>A200293 001</u>	Nov 09, 2021
AN	WILSHIRE PHARMS INC	<u>EQ 0.015MG BASE/2ML</u>	<u>A216815 001</u>	Nov 25, 2022

BROVANA**AN** +! LUPIN **EQ 0.015MG BASE/2ML** **N021912 001** Oct 06, 2006ARGATROBAN

INJECTABLE; INJECTION

ARGATROBAN

AP	AMNEAL PHARMS CO	<u>250MG/2.5ML (100MG/ML)</u>	<u>A206698 001</u>	Jan 26, 2018
AP	CAPLIN	<u>50MG/50ML (1MG/ML)</u>	<u>A214235 001</u>	Jan 21, 2021
AP	ENDO OPERATIONS	<u>250MG/2.5ML (100MG/ML)</u>	<u>A091665 001</u>	Jun 30, 2014
AP	FRESENIUS KABI USA	<u>250MG/2.5ML (100MG/ML)</u>	<u>N201811 001</u>	Mar 23, 2015
AP	GLAND PHARMA LTD	<u>50MG/50ML (1MG/ML)</u>	<u>A217848 001</u>	Jul 31, 2023
AP	+! HIKMA PHARM CO LTD	<u>50MG/50ML (1MG/ML)</u>	<u>N203049 002</u>	Sep 30, 2016
AP	+! HOSPIRA	<u>250MG/2.5ML (100MG/ML)</u>	<u>N203049 001</u>	Jan 05, 2012
AP	! HOSPIRA	<u>250MG/2.5ML (100MG/ML)</u>	<u>A204120 001</u>	Sep 21, 2016
AP	MYLAN INSTITUTIONAL	<u>250MG/2.5ML (100MG/ML)</u>	<u>A202626 001</u>	Jun 30, 2014

INJECTABLE; INTRAVENOUS

ARGATROBAN IN SODIUM CHLORIDE

AP	GLAND PHARMA LTD	<u>125MG/125ML (1MG/ML)</u>	<u>A205570 001</u>	May 22, 2017
AP	+! SANDOZ	<u>125MG/125ML (1MG/ML)</u>	<u>N022485 001</u>	May 09, 2011

SOLUTION; INTRAVENOUS

ARGATROBAN IN SODIUM CHLORIDE

+! ACCORD HLTHCARE 50MG/50ML (1MG/ML) N212035 001 Jun 07, 2021

EUGIA PHARMA 50MG/50ML (1MG/ML) N209552 001 Nov 27, 2018

SPECLTS

ARGININE HYDROCHLORIDE

INJECTABLE; INJECTION

R-GENE 10

+! PHARMACIA AND UPJOHN 10GM/100ML N016931 001

ARIMOCLOMOL CITRATE

CAPSULE; ORAL

MIPLYFFA

+ ZEVRA DENMARK EQ 47MG BASE N214927 001 Sep 20, 2024

+ EQ 62MG BASE N214927 002 Sep 20, 2024

+ EQ 93MG BASE N214927 003 Sep 20, 2024

+! EQ 124MG BASE N214927 004 Sep 20, 2024

PRESCRIPTION DRUG PRODUCT LIST

ARIPIPIRAZOLE

FILM;ORAL

OPIPZA

+	XIAMEN LP PHARM CO	2MG	N216655	001	Jul 22, 2024
+		5MG	N216655	002	Jul 22, 2024
+	!	10MG	N216655	003	Jul 22, 2024

FOR SUSPENSION, EXTENDED RELEASE;INTRAMUSCULAR

ABILIFY MAINTENA KIT

AP	+	OTSUKA PHARM CO LTD	300MG/VIAL	N202971	001	Feb 28, 2013
AP	+	!	400MG/VIAL	N202971	002	Feb 28, 2013

ARIPIPIRAZOLE

AP		MYLAN	300MG/VIAL	A216608	001	Dec 03, 2024
AP			400MG/VIAL	A216608	002	Dec 03, 2024

ABILIFY MAINTENA KIT

+	OTSUKA PHARM CO LTD	300MG	N202971	003	Sep 29, 2014
+		400MG	N202971	004	Sep 29, 2014

SOLUTION;ORAL

ARIPIPIRAZOLE

AA	!	AMNEAL PHARMS	1MG/ML	A203906	001	Aug 14, 2015
AA		APOTEX	1MG/ML	A204094	001	Sep 30, 2015
AA		AUROBINDO PHARMA LTD	1MG/ML	A210479	001	Jan 29, 2019
AA		CHARTWELL RX	1MG/ML	A215595	001	Oct 25, 2022
AA		HETERO LABS LTD III	1MG/ML	A216150	001	Nov 01, 2023
AA		LANNETT CO INC	1MG/ML	A204171	001	Aug 14, 2015
AA		RUBICON	1MG/ML	A216351	001	Oct 27, 2022
AA		VISTAPHARM	1MG/ML	A212870	001	Dec 26, 2019

SUSPENSION, EXTENDED RELEASE;INTRAMUSCULAR

ABILIFY ASIMTUFI

+	OTSUKA	720MG/2.4ML (300MG/ML)	N217006	001	Apr 27, 2023
+	!	960MG/3.2ML (300MG/ML)	N217006	002	Apr 27, 2023

TABLET;ORAL

ABILIFY

AB	+	OTSUKA	2MG	N021436	006	Nov 15, 2002
AB	+		5MG	N021436	005	Nov 15, 2002
AB	+	!	10MG	N021436	001	Nov 15, 2002
AB	+		15MG	N021436	002	Nov 15, 2002
AB	+		20MG	N021436	003	Nov 15, 2002
AB	+		30MG	N021436	004	Nov 15, 2002

ARIPIPIRAZOLE

AB		ACCORD HLTHCARE	2MG	A206251	001	Dec 07, 2016
AB			5MG	A206251	002	Dec 07, 2016
AB			10MG	A206251	003	Dec 07, 2016
AB			15MG	A206251	004	Dec 07, 2016
AB			20MG	A206251	005	Dec 07, 2016
AB			30MG	A206251	006	Dec 07, 2016
AB		AJANTA PHARMA LTD	2MG	A206174	001	Sep 12, 2016
AB			5MG	A206174	002	Sep 12, 2016
AB			10MG	A206174	003	Sep 12, 2016
AB			15MG	A206174	004	Sep 12, 2016
AB			20MG	A206174	005	Sep 12, 2016
AB			30MG	A206174	006	Sep 12, 2016
AB		ALEMbic	2MG	A202101	001	Apr 28, 2015
AB			5MG	A202101	002	Apr 28, 2015
AB			10MG	A202101	003	Apr 28, 2015
AB			15MG	A202101	004	Apr 28, 2015
AB			20MG	A202101	005	Apr 28, 2015
AB			30MG	A202101	006	Apr 28, 2015
AB		ALKEM LABS LTD	2MG	A207105	001	Feb 21, 2019
AB			5MG	A207105	002	Feb 21, 2019
AB			10MG	A207105	003	Feb 21, 2019
AB			15MG	A207105	004	Feb 21, 2019
AB			20MG	A207105	005	Feb 21, 2019
AB			30MG	A207105	006	Feb 21, 2019
AB		AMNEAL PHARMS	2MG	A204838	001	Jun 17, 2016
AB			5MG	A204838	002	Jun 17, 2016
AB			10MG	A204838	003	Jun 17, 2016
AB			15MG	A204838	004	Jun 17, 2016
AB			20MG	A204838	005	Jun 17, 2016
AB			30MG	A204838	006	Jun 17, 2016
AB		APOTEX	2MG	A078583	001	Jul 24, 2015
AB			5MG	A078583	002	Jul 24, 2015
AB			10MG	A078583	003	Jul 24, 2015

PRESCRIPTION DRUG PRODUCT LIST

ARIPIPIRAZOLE

TABLET; ORAL

ARIPIPIRAZOLE

<u>AB</u>		<u>15MG</u>	<u>A078583</u>	<u>004</u>	Jul 24, 2015
<u>AB</u>		<u>20MG</u>	<u>A078583</u>	<u>005</u>	Jul 24, 2015
<u>AB</u>		<u>30MG</u>	<u>A078583</u>	<u>006</u>	Jul 24, 2015
<u>AB</u>	AUROBINDO PHARMA	<u>2MG</u>	<u>A203908</u>	<u>001</u>	Oct 08, 2015
<u>AB</u>		<u>5MG</u>	<u>A203908</u>	<u>002</u>	Oct 08, 2015
<u>AB</u>		<u>10MG</u>	<u>A203908</u>	<u>003</u>	Oct 08, 2015
<u>AB</u>		<u>15MG</u>	<u>A203908</u>	<u>004</u>	Oct 08, 2015
<u>AB</u>		<u>20MG</u>	<u>A203908</u>	<u>005</u>	Oct 08, 2015
<u>AB</u>		<u>30MG</u>	<u>A203908</u>	<u>006</u>	Oct 08, 2015
<u>AB</u>	HETERO LABS LTD V	<u>2MG</u>	<u>A205064</u>	<u>001</u>	Apr 28, 2015
<u>AB</u>		<u>5MG</u>	<u>A205064</u>	<u>002</u>	Apr 28, 2015
<u>AB</u>		<u>10MG</u>	<u>A205064</u>	<u>003</u>	Apr 28, 2015
<u>AB</u>		<u>15MG</u>	<u>A205064</u>	<u>004</u>	Apr 28, 2015
<u>AB</u>		<u>20MG</u>	<u>A205064</u>	<u>005</u>	Apr 28, 2015
<u>AB</u>		<u>30MG</u>	<u>A205064</u>	<u>006</u>	Apr 28, 2015
<u>AB</u>	LUPIN	<u>2MG</u>	<u>A205589</u>	<u>001</u>	Mar 18, 2024
<u>AB</u>		<u>5MG</u>	<u>A205589</u>	<u>002</u>	Mar 18, 2024
<u>AB</u>		<u>10MG</u>	<u>A205589</u>	<u>003</u>	Mar 18, 2024
<u>AB</u>		<u>15MG</u>	<u>A205589</u>	<u>004</u>	Mar 18, 2024
<u>AB</u>		<u>20MG</u>	<u>A205589</u>	<u>005</u>	Mar 18, 2024
<u>AB</u>		<u>30MG</u>	<u>A205589</u>	<u>006</u>	Mar 18, 2024
<u>AB</u>	MACLEODS PHARMS LTD	<u>2MG</u>	<u>A204111</u>	<u>001</u>	Oct 07, 2016
<u>AB</u>		<u>5MG</u>	<u>A204111</u>	<u>002</u>	Oct 07, 2016
<u>AB</u>		<u>10MG</u>	<u>A204111</u>	<u>003</u>	Oct 07, 2016
<u>AB</u>		<u>15MG</u>	<u>A204111</u>	<u>004</u>	Oct 07, 2016
<u>AB</u>		<u>20MG</u>	<u>A204111</u>	<u>005</u>	Oct 07, 2016
<u>AB</u>		<u>30MG</u>	<u>A204111</u>	<u>006</u>	Oct 07, 2016
<u>AB</u>	ORBION PHARMS	<u>2MG</u>	<u>A202683</u>	<u>001</u>	May 23, 2017
<u>AB</u>		<u>5MG</u>	<u>A202683</u>	<u>002</u>	May 23, 2017
<u>AB</u>		<u>10MG</u>	<u>A202683</u>	<u>003</u>	May 23, 2017
<u>AB</u>		<u>15MG</u>	<u>A202683</u>	<u>004</u>	May 23, 2017
<u>AB</u>		<u>20MG</u>	<u>A202683</u>	<u>005</u>	May 23, 2017
<u>AB</u>		<u>30MG</u>	<u>A202683</u>	<u>006</u>	May 23, 2017
<u>AB</u>	PRINSTON INC	<u>2MG</u>	<u>A205363</u>	<u>001</u>	Dec 04, 2017
<u>AB</u>		<u>5MG</u>	<u>A205363</u>	<u>002</u>	Dec 04, 2017
<u>AB</u>		<u>10MG</u>	<u>A205363</u>	<u>003</u>	Dec 04, 2017
<u>AB</u>		<u>15MG</u>	<u>A205363</u>	<u>004</u>	Dec 04, 2017
<u>AB</u>		<u>20MG</u>	<u>A205363</u>	<u>005</u>	Dec 04, 2017
<u>AB</u>		<u>30MG</u>	<u>A205363</u>	<u>006</u>	Dec 04, 2017
<u>AB</u>	SCIEGEN PHARMS INC	<u>2MG</u>	<u>A206383</u>	<u>001</u>	Sep 29, 2016
<u>AB</u>		<u>5MG</u>	<u>A206383</u>	<u>002</u>	Sep 29, 2016
<u>AB</u>		<u>10MG</u>	<u>A206383</u>	<u>003</u>	Sep 29, 2016
<u>AB</u>		<u>15MG</u>	<u>A206383</u>	<u>004</u>	Sep 29, 2016
<u>AB</u>		<u>20MG</u>	<u>A206383</u>	<u>005</u>	Sep 29, 2016
<u>AB</u>		<u>30MG</u>	<u>A206383</u>	<u>006</u>	Sep 29, 2016
<u>AB</u>	SUNSHINE	<u>2MG</u>	<u>A213037</u>	<u>001</u>	Oct 02, 2023
<u>AB</u>		<u>5MG</u>	<u>A213037</u>	<u>002</u>	Oct 02, 2023
<u>AB</u>		<u>10MG</u>	<u>A213037</u>	<u>003</u>	Oct 02, 2023
<u>AB</u>	TORRENT	<u>2MG</u>	<u>A201519</u>	<u>001</u>	Apr 28, 2015
<u>AB</u>		<u>10MG</u>	<u>A201519</u>	<u>003</u>	Apr 28, 2015
<u>AB</u>		<u>5MG</u>	<u>A201519</u>	<u>002</u>	Apr 28, 2015
<u>AB</u>		<u>15MG</u>	<u>A201519</u>	<u>004</u>	Apr 28, 2015
<u>AB</u>		<u>20MG</u>	<u>A201519</u>	<u>005</u>	Apr 28, 2015
<u>AB</u>		<u>30MG</u>	<u>A201519</u>	<u>006</u>	Apr 28, 2015
<u>AB</u>	UNICHEM	<u>2MG</u>	<u>A203025</u>	<u>001</u>	Dec 01, 2021
<u>AB</u>		<u>5MG</u>	<u>A203025</u>	<u>002</u>	Dec 01, 2021
<u>AB</u>		<u>10MG</u>	<u>A203025</u>	<u>003</u>	Dec 01, 2021
<u>AB</u>		<u>15MG</u>	<u>A203025</u>	<u>004</u>	Dec 01, 2021
<u>AB</u>		<u>20MG</u>	<u>A203025</u>	<u>005</u>	Dec 01, 2021
<u>AB</u>		<u>30MG</u>	<u>A203025</u>	<u>006</u>	Dec 01, 2021

TABLET, ORALLY DISINTEGRATING; ORAL

ARIPIPIRAZOLE

<u>AB</u>	! ALEMBIC	<u>10MG</u>	<u>A202102</u>	<u>001</u>	Apr 28, 2015
<u>AB</u>		<u>15MG</u>	<u>A202102</u>	<u>002</u>	Apr 28, 2015
<u>AB</u>	ORBION PHARMS	<u>10MG</u>	<u>A202547</u>	<u>001</u>	Dec 11, 2017
<u>AB</u>		<u>15MG</u>	<u>A202547</u>	<u>002</u>	Dec 11, 2017
<u>AB</u>	SCIEGEN PHARMS INC	<u>10MG</u>	<u>A207240</u>	<u>001</u>	Apr 18, 2018
<u>AB</u>		<u>15MG</u>	<u>A207240</u>	<u>002</u>	Apr 18, 2018
<u>AB</u>	SQUARE PHARMS	<u>10MG</u>	<u>A090165</u>	<u>001</u>	Aug 28, 2018

PRESCRIPTION DRUG PRODUCT LIST

ARIPIRAZOLE

TABLET, ORALLY DISINTEGRATING;ORAL

ARIPIRAZOLE

<u>AB</u>		<u>15MG</u>	<u>A090165 002</u>	Aug 28, 2018
		20MG	A090165 003	Aug 28, 2018
		30MG	A090165 004	Aug 28, 2018

ARIPIRAZOLE LAUROXIL

SUSPENSION, EXTENDED RELEASE;INTRAMUSCULAR

ARISTADA

+	ALKERMES INC	441MG/1.6ML (275.63MG/ML)	N207533 001	Oct 05, 2015	
+		662MG/2.4ML (275.83MG/ML)	N207533 002	Oct 05, 2015	
+	!	882MG/3.2ML (275.63MG/ML)	N207533 003	Oct 05, 2015	
+		1064MG/3.9ML (272.82MG/ML)	N207533 004	Jun 05, 2017	
	ARISTADA INITIO KIT				
+	!	ALKERMES INC	675MG/2.4ML (281.25MG/ML)	N209830 001	Jun 29, 2018

ARMODAFINIL

TABLET;ORAL

ARMODAFINIL

<u>AB</u>	AUROBINDO PHARMA LTD	<u>50MG</u>	<u>A206069 001</u>	Mar 06, 2018	
<u>AB</u>		<u>150MG</u>	<u>A206069 002</u>	Mar 06, 2018	
<u>AB</u>		<u>200MG</u>	<u>A206069 004</u>	Dec 07, 2018	
<u>AB</u>		<u>250MG</u>	<u>A206069 003</u>	Mar 06, 2018	
<u>AB</u>	LUPIN LTD	<u>50MG</u>	<u>A200751 001</u>	Nov 28, 2016	
<u>AB</u>		<u>150MG</u>	<u>A200751 003</u>	Nov 28, 2016	
<u>AB</u>		<u>200MG</u>	<u>A200751 004</u>	Nov 28, 2016	
<u>AB</u>		<u>250MG</u>	<u>A200751 005</u>	Nov 28, 2016	
<u>AB</u>	MYLAN PHARMS INC	<u>50MG</u>	<u>A200043 001</u>	Jun 01, 2012	
<u>AB</u>		<u>100MG</u>	<u>A200043 004</u>	May 09, 2019	
<u>AB</u>		<u>150MG</u>	<u>A200043 002</u>	Jun 01, 2012	
<u>AB</u>		<u>200MG</u>	<u>A200043 005</u>	May 09, 2019	
<u>AB</u>		<u>250MG</u>	<u>A200043 003</u>	Jun 01, 2012	
<u>AB</u>	NATCO PHARMA LTD	<u>50MG</u>	<u>A202768 001</u>	Nov 28, 2016	
<u>AB</u>		<u>100MG</u>	<u>A202768 004</u>	Sep 28, 2017	
<u>AB</u>		<u>150MG</u>	<u>A202768 002</u>	Nov 28, 2016	
<u>AB</u>		<u>200MG</u>	<u>A202768 005</u>	Sep 28, 2017	
<u>AB</u>		<u>250MG</u>	<u>A202768 003</u>	Nov 28, 2016	
	<u>NUVIGIL</u>				
<u>AB</u>	+	CEPHALON	<u>50MG</u>	<u>N021875 001</u>	Jun 15, 2007
<u>AB</u>	+		<u>150MG</u>	<u>N021875 003</u>	Jun 15, 2007
<u>AB</u>	+		<u>200MG</u>	<u>N021875 005</u>	Mar 26, 2009
<u>AB</u>	+	!	<u>250MG</u>	<u>N021875 004</u>	Jun 15, 2007

ARSENIC TRIOXIDE

INJECTABLE; INJECTION

ARSENIC TRIOXIDE

<u>AP</u>	AMNEAL	<u>1MG/ML</u>	<u>A210739 001</u>	Jan 25, 2021	
<u>AP</u>		<u>2MG/ML</u>	<u>A210739 002</u>	Aug 19, 2021	
<u>AP</u>	AMRING PHARMS	<u>1MG/ML</u>	<u>A210802 001</u>	Nov 13, 2018	
<u>AP</u>	!	FRESENIUS KABI USA	<u>1MG/ML</u>	<u>A208231 001</u>	Aug 31, 2018
<u>AP</u>	GLAND PHARMA LTD	<u>1MG/ML</u>	<u>A215059 002</u>	Apr 22, 2022	
<u>AP</u>		<u>2MG/ML</u>	<u>A215059 001</u>	Oct 07, 2021	
<u>AP</u>	INGENUS PHARMS LLC	<u>1MG/ML</u>	<u>A209315 001</u>	Nov 15, 2018	
<u>AP</u>	NEXUS	<u>1MG/ML</u>	<u>A209780 001</u>	Nov 15, 2018	
<u>AP</u>	ORBICULAR	<u>1MG/ML</u>	<u>A217413 001</u>	Apr 20, 2023	
<u>AP</u>		<u>2MG/ML</u>	<u>A217413 002</u>	Apr 20, 2023	
<u>AP</u>	PENN LIFE	<u>1MG/ML</u>	<u>A209873 001</u>	May 06, 2019	
<u>AP</u>	SANDOZ	<u>2MG/ML</u>	<u>A215359 001</u>	Dec 02, 2021	
<u>AP</u>	ZYDUS PHARMS	<u>1MG/ML</u>	<u>A206228 001</u>	Nov 13, 2018	
<u>AP</u>		<u>2MG/ML</u>	<u>A206228 002</u>	Aug 30, 2019	

TRISENOX

<u>AP</u>	+	!	CEPHALON	<u>2MG/ML</u>	<u>N021248 002</u>	Oct 13, 2017
-----------	---	---	----------	---------------	--------------------	--------------

ARTEMETHER; LUMEFANTRINE

TABLET;ORAL

COARTEM

+	!	NOVARTIS	20MG;120MG	N022268 001	Apr 07, 2009
---	---	----------	------------	-------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

ARTESUNATE

POWDER; INTRAVENOUS

ARTESUNATE

+! AMIVAS 110MG/VIAL N213036 001 May 26, 2020

ARTICAINE HYDROCHLORIDE; EPINEPHRINE BITARTRATE

INJECTABLE; INJECTION

ORABLOC

+ PIERREL 4%;EQ 0.009MG BASE/1.8ML (EQ 0.005MG BASE/ML) N022466 001 Feb 26, 2010

+! 4%;EQ 0.018MG BASE/1.8ML (EQ 0.01MG BASE/ML) N022466 002 Feb 26, 2010

SEPTOCAINE

+! DEPROCO 4%; EQ 0.0085MG BASE/1.7ML (4%; EQ 0.005MG BASE/ML) N020971 002 Mar 30, 2006

+! 4%;EQ 0.017MG BASE/1.7ML (4%;EQ 0.01MG BASE/ML) N020971 001 Apr 03, 2000

ASCIMINIB HYDROCHLORIDE

TABLET; ORAL

SCEMBLIX

+ NOVARTIS EQ 20MG BASE N215358 001 Oct 29, 2021

+! EQ 40MG BASE N215358 002 Oct 29, 2021

+ EQ 100MG BASE N215358 003 Apr 18, 2024

ASCORBIC ACID

SOLUTION; INTRAVENOUS

ASCOR

+! MCGUFF 25,000MG/50ML (500MG/ML) N209112 001 Oct 02, 2017

ASCORBIC ACID; BIOTIN; CHOLECALCIFEROL; CYANOCOBALAMIN; DEXPANTHENOL; FOLIC ACID; NIACINAMIDE; PYRIDOXINE; RIBOFLAVIN; THIAMINE; TOCOPHEROL ACETATE; VITAMIN A; VITAMIN K

INJECTABLE; INTRAVENOUS

INFUVITE PEDIATRIC

+! SANDOZ CANADA INC 80MG/VIAL;0.02MG/VIAL;400 N021265 001 Feb 21, 2001

IU/VIAL;0.001MG/VIAL;5MG/VIAL;0.14MG/VIAL;17MG/VIAL;1MG/VIAL;1.4MG/VIAL;1.2MG/VIAL;7 IU/VIAL;2,300 IU/VIAL;0.2MG/VIAL

INFUVITE PEDIATRIC (PHARMACY BULK PACKAGE)

+! SANDOZ CANADA INC 80MG/VIAL;0.02MG/VIAL;400 N021265 002 Jan 29, 2004

IU/VIAL;0.001MG/VIAL;5MG/VIAL;0.14MG/VIAL;17MG/VIAL;1MG/VIAL;1.4MG/VIAL;1.2MG/VIAL;7 IU/VIAL;2,300 IU/VIAL;0.2MG/VIAL

ASCORBIC ACID; POLYETHYLENE GLYCOL 3350; POTASSIUM CHLORIDE; SODIUM ASCORBATE; SODIUM CHLORIDE; SODIUM SULFATE

FOR SOLUTION; ORAL

MOVIPREPAA +! SALIX PHARMS 4.7GM;100GM;1.015GM;5.9GM;2.691GM;7.5GM N021881 001 Aug 02, 2006PEG-3350, SODIUM SULFATE, SODIUM CHLORIDE, POTASSIUM CHLORIDE, SODIUM ASCORBATE AND ASCORBICAA NOVEL LABS INC 4.7GM;100GM;1.015GM;5.9GM;2.691GM;7.5GM A090145 001 Jan 25, 2012AA TARO 4.7GM;100GM;1.015GM;5.9GM;2.691GM;7.5GM A207498 001 Jun 28, 2024

PLENVU

+! SALIX 7.54GM;140GM;2.2GM;48.11GM;5.2GM;9GM N209381 001 May 04, 2018

ASENAPINE

SYSTEM; TRANSDERMAL

SECUADO

+! HISAMITSU 3.8MG/24HR N212268 001 Oct 11, 2019

+ 5.7MG/24HR N212268 002 Oct 11, 2019

+ 7.6MG/24HR N212268 003 Oct 11, 2019

ASENAPINE MALEATE

TABLET; SUBLINGUAL

ASENAPINE MALEATEAB ALEMBIC EQ 2.5MG BASE A206098 003 Jul 19, 2021AB EQ 5MG BASE A206098 001 Dec 10, 2020AB EQ 10MG BASE A206098 002 Dec 10, 2020AB BRECKENRIDGE EQ 2.5MG BASE A205960 001 Dec 10, 2020AB EQ 5MG BASE A205960 003 Mar 08, 2021AB EQ 10MG BASE A205960 002 Dec 10, 2020AB SIGMAPHARM LABS LLC EQ 5MG BASE A206107 001 Dec 10, 2020AB EQ 10MG BASE A206107 002 Dec 10, 2020SAPHRISAB + ALLERGAN EQ 2.5MG BASE N022117 003 Mar 12, 2015AB + EQ 5MG BASE N022117 001 Aug 13, 2009AB +! EQ 10MG BASE N022117 002 Aug 13, 2009

PRESCRIPTION DRUG PRODUCT LIST

ASPIRIN; BUTALBITAL; CAFFEINE

CAPSULE; ORAL

BUTALBITAL, ASPIRIN AND CAFFEINE

AA	NOSTRUM LABS INC	325MG; 50MG; 40MG	A078149 001	Jun 13, 2007
-----------	------------------	--------------------------	--------------------	--------------

LANORINAL

AA	! SANDOZ	325MG; 50MG; 40MG	A086996 002	Oct 11, 1985
-----------	----------	--------------------------	--------------------	--------------

TABLET; ORAL

BUTALBITAL, ASPIRIN AND CAFFEINE

!	STRIDES PHARMA	325MG; 50MG; 40MG	A204195 001	Sep 22, 2016
---	----------------	-------------------	-------------	--------------

ASPIRIN; BUTALBITAL; CAFFEINE; CODEINE PHOSPHATE

CAPSULE; ORAL

BUTALBITAL, ASPIRIN, CAFFEINE, AND CODEINE PHOSPHATE

AB	LGM PHARMA	325MG; 50MG; 40MG; 30MG	A075231 001	Nov 30, 2001
-----------	------------	--------------------------------	--------------------	--------------

AB	! STEVENS J	325MG; 50MG; 40MG; 30MG	A074951 001	Aug 31, 1998
-----------	-------------	--------------------------------	--------------------	--------------

ASPIRIN; CAFFEINE; ORPHENADRINE CITRATE

TABLET; ORAL

ORPHENGESIC

	GALT PHARMS	385MG; 30MG; 25MG	A075141 001	May 29, 1998
--	-------------	-------------------	-------------	--------------

ORPHENGESIC FORTE

!	GALT PHARMS	770MG; 60MG; 50MG	A075141 002	May 29, 1998
---	-------------	-------------------	-------------	--------------

ASPIRIN; DIPYRIDAMOLE

CAPSULE, EXTENDED RELEASE; ORAL

ASPIRIN AND DIPYRIDAMOLE

AB	AMNEAL PHARMS	25MG; 200MG	A206392 001	Mar 08, 2016
-----------	---------------	--------------------	--------------------	--------------

AB	BARR	25MG; 200MG	A078804 001	Aug 14, 2009
-----------	------	--------------------	--------------------	--------------

AB	DR REDDYS	25MG; 200MG	A209048 001	Oct 10, 2018
-----------	-----------	--------------------	--------------------	--------------

AB	ENDO OPERATIONS	25MG; 200MG	A207944 001	Jan 18, 2017
-----------	-----------------	--------------------	--------------------	--------------

AB	! GLENMARK SPECLT	25MG; 200MG	A210318 001	May 24, 2019
-----------	-------------------	--------------------	--------------------	--------------

AB	MICRO LABS	25MG; 200MG	A209929 001	Aug 11, 2021
-----------	------------	--------------------	--------------------	--------------

AB	SANDOZ	25MG; 200MG	A206739 001	Jan 18, 2017
-----------	--------	--------------------	--------------------	--------------

AB	ZYDUS PHARMS	25MG; 200MG	A206753 001	Aug 29, 2017
-----------	--------------	--------------------	--------------------	--------------

ASPIRIN; HYDROCODONE BITARTRATE

TABLET; ORAL

HYDROCODONE BITARTRATE AND ASPIRIN

!	LGM PHARMA	500MG; 5MG	A205479 001	May 28, 2021
---	------------	------------	-------------	--------------

ASPIRIN; OXYCODONE HYDROCHLORIDE

TABLET; ORAL

OXYCODONE AND ASPIRIN

AA	EPIC PHARMA LLC	325MG; 4.8355MG	A040910 001	Jul 16, 2020
-----------	-----------------	------------------------	--------------------	--------------

PERCODAN

AA	+! ENDO OPERATIONS	325MG; 4.8355MG	N007337 007	Aug 05, 2005
-----------	--------------------	------------------------	--------------------	--------------

ATAZANAVIR SULFATE

CAPSULE; ORAL

ATAZANAVIR SULFATE

AB	AUROBINDO PHARMA	EQ 100MG BASE	A204806 001	Jun 25, 2018
-----------	------------------	----------------------	--------------------	--------------

AB		EQ 150MG BASE	A204806 002	Jun 25, 2018
-----------	--	----------------------	--------------------	--------------

AB		EQ 200MG BASE	A204806 003	Jun 25, 2018
-----------	--	----------------------	--------------------	--------------

AB		EQ 300MG BASE	A204806 004	Jun 25, 2018
-----------	--	----------------------	--------------------	--------------

AB	HETERO LABS LTD III	EQ 150MG BASE	A212278 001	Feb 02, 2022
-----------	---------------------	----------------------	--------------------	--------------

AB		EQ 200MG BASE	A212278 002	Feb 02, 2022
-----------	--	----------------------	--------------------	--------------

AB		EQ 300MG BASE	A212278 003	Feb 02, 2022
-----------	--	----------------------	--------------------	--------------

AB	LAURUS	EQ 150MG BASE	A212579 001	Apr 30, 2021
-----------	--------	----------------------	--------------------	--------------

AB		EQ 200MG BASE	A212579 002	Apr 30, 2021
-----------	--	----------------------	--------------------	--------------

AB		EQ 300MG BASE	A212579 003	Apr 30, 2021
-----------	--	----------------------	--------------------	--------------

AB	TEVA PHARMS USA	EQ 100MG BASE	A091673 001	Apr 22, 2014
-----------	-----------------	----------------------	--------------------	--------------

AB		EQ 150MG BASE	A091673 002	Apr 22, 2014
-----------	--	----------------------	--------------------	--------------

AB		EQ 200MG BASE	A091673 003	Apr 22, 2014
-----------	--	----------------------	--------------------	--------------

AB		EQ 300MG BASE	A091673 004	Apr 22, 2014
-----------	--	----------------------	--------------------	--------------

REYATAZ

AB	+ BRISTOL MYERS	EQ 200MG BASE	N021567 003	Jun 20, 2003
-----------	-----------------	----------------------	--------------------	--------------

AB	+! SQUIBB	EQ 300MG BASE	N021567 004	Oct 16, 2006
-----------	-----------	----------------------	--------------------	--------------

POWDER; ORAL

REYATAZ

+	BRISTOL MYERS	EQ 50MG BASE/PACKET	N206352 001	Jun 02, 2014
---	---------------	---------------------	-------------	--------------

SQUIBB

PRESCRIPTION DRUG PRODUCT LIST

ATAZANAVIR SULFATE; COBICISTAT

TABLET; ORAL

EVOTAZ

+! BRISTOL

EQ 300MG BASE;150MG

N206353 001 Jan 29, 2015

ATENOLOL

TABLET; ORAL

ATENOLOL

<u>AB</u>	AUROBINDO PHARMA	<u>25MG</u>	<u>A078512 001</u>	Oct 31, 2007
<u>AB</u>		<u>50MG</u>	<u>A078512 002</u>	Oct 31, 2007
<u>AB</u>		<u>100MG</u>	<u>A078512 003</u>	Oct 31, 2007
<u>AB</u>	HLTHCARE	<u>25MG</u>	<u>A073026 002</u>	May 01, 1992
<u>AB</u>		<u>50MG</u>	<u>A073026 003</u>	Sep 17, 1991
<u>AB</u>		<u>100MG</u>	<u>A073026 001</u>	Sep 17, 1991
<u>AB</u>	MYLAN	<u>25MG</u>	<u>A073457 002</u>	Apr 26, 1999
<u>AB</u>		<u>50MG</u>	<u>A073457 003</u>	Jan 24, 1992
<u>AB</u>		<u>100MG</u>	<u>A073457 001</u>	Jan 24, 1992
<u>AB</u>	TEVA	<u>25MG</u>	<u>A074056 003</u>	Jul 19, 2004
<u>AB</u>		<u>50MG</u>	<u>A074056 001</u>	Jan 18, 1995
<u>AB</u>		<u>100MG</u>	<u>A074056 002</u>	Jan 18, 1995
<u>AB</u>	TWI PHARMS	<u>25MG</u>	<u>A072304 002</u>	Jul 31, 1992
<u>AB</u>		<u>50MG</u>	<u>A072304 003</u>	Jul 18, 1988
<u>AB</u>		<u>100MG</u>	<u>A072304 001</u>	Jul 15, 1988
<u>AB</u>	UNICHEM	<u>25MG</u>	<u>A213136 001</u>	Nov 21, 2019
<u>AB</u>		<u>50MG</u>	<u>A213136 002</u>	Nov 21, 2019
<u>AB</u>		<u>100MG</u>	<u>A213136 003</u>	Nov 21, 2019
<u>AB</u>	UNIQUE	<u>25MG</u>	<u>A077443 001</u>	Sep 13, 2006
<u>AB</u>		<u>50MG</u>	<u>A077443 002</u>	Sep 13, 2006
<u>AB</u>		<u>100MG</u>	<u>A077443 003</u>	Sep 13, 2006
<u>AB</u>	ZYDUS PHARMS USA	<u>25MG</u>	<u>A076900 001</u>	Jan 28, 2005
<u>AB</u>		<u>50MG</u>	<u>A076900 002</u>	Jan 28, 2005
<u>AB</u>		<u>100MG</u>	<u>A076900 003</u>	Jan 28, 2005

TENORMIN

<u>AB</u>	+ TWI PHARMS	<u>25MG</u>	<u>N018240 004</u>	Apr 09, 1990
<u>AB</u>	+	<u>50MG</u>	<u>N018240 001</u>	
<u>AB</u>	+!	<u>100MG</u>	<u>N018240 002</u>	

ATENOLOL; CHLORTHALIDONE

TABLET; ORAL

ATENOLOL AND CHLORTHALIDONE

<u>AB</u>	NOVITIUM PHARMA	<u>50MG;25MG</u>	<u>A215560 001</u>	Oct 25, 2021
<u>AB</u>		<u>100MG;25MG</u>	<u>A215560 002</u>	Oct 25, 2021
<u>AB</u>	TWI PHARMS	<u>50MG;25MG</u>	<u>A072302 002</u>	May 31, 1990
<u>AB</u>		<u>100MG;25MG</u>	<u>A072302 001</u>	May 31, 1990
<u>AB</u>	UNICHEM	<u>50MG;25MG</u>	<u>A213302 001</u>	Nov 25, 2020
<u>AB</u>		<u>100MG;25MG</u>	<u>A213302 002</u>	Nov 25, 2020
<u>AB</u>	WATSON LABS	<u>50MG;25MG</u>	<u>A073665 001</u>	Jul 02, 1992
<u>AB</u>		<u>100MG;25MG</u>	<u>A073665 002</u>	Jul 02, 1992
<u>AB</u>	ZYDUS PHARMS	<u>50MG;25MG</u>	<u>A210028 001</u>	Mar 08, 2019
<u>AB</u>		<u>100MG;25MG</u>	<u>A210028 002</u>	Mar 08, 2019

TENORETIC 100

<u>AB</u>	+! TWI PHARMS	<u>100MG;25MG</u>	<u>N018760 001</u>	Jun 08, 1984
-----------	---------------	-------------------	--------------------	--------------

TENORETIC 50

<u>AB</u>	+ TWI PHARMS	<u>50MG;25MG</u>	<u>N018760 002</u>	Jun 08, 1984
-----------	--------------	------------------	--------------------	--------------

ATOGEFANT

TABLET; ORAL

QULIPTA

+ ABBVIE

10MG

N215206 001 Sep 28, 2021

+

30MG

N215206 002 Sep 28, 2021

+!

60MG

N215206 003 Sep 28, 2021

ATOMOXETINE HYDROCHLORIDE

CAPSULE; ORAL

ATOMOXETINE HYDROCHLORIDE

<u>AB</u>	APOTEX	<u>EQ 10MG BASE</u>	<u>A078983 001</u>	May 30, 2017
<u>AB</u>		<u>EQ 18MG BASE</u>	<u>A078983 002</u>	May 30, 2017
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>A078983 003</u>	May 30, 2017
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A078983 004</u>	May 30, 2017
<u>AB</u>		<u>EQ 60MG BASE</u>	<u>A078983 005</u>	May 30, 2017
<u>AB</u>		<u>EQ 80MG BASE</u>	<u>A078983 006</u>	May 30, 2017
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A078983 007</u>	May 30, 2017
<u>AB</u>	AUROBINDO PHARMA	<u>EQ 10MG BASE</u>	<u>A079016 001</u>	May 30, 2017
<u>AB</u>		<u>EQ 18MG BASE</u>	<u>A079016 002</u>	May 30, 2017

PRESCRIPTION DRUG PRODUCT LIST

ATOMOXETINE HYDROCHLORIDE

CAPSULE; ORAL

ATOMOXETINE HYDROCHLORIDE

<u>AB</u>		<u>EQ 25MG BASE</u>	<u>A079016 003</u>	May 30, 2017
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A079016 004</u>	May 30, 2017
<u>AB</u>		<u>EQ 60MG BASE</u>	<u>A079016 005</u>	May 30, 2017
<u>AB</u>		<u>EQ 80MG BASE</u>	<u>A079016 006</u>	May 30, 2017
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A079016 007</u>	May 30, 2017
<u>AB</u>	DR REDDYS	<u>EQ 10MG BASE</u>	<u>A090609 001</u>	Feb 23, 2018
<u>AB</u>		<u>EQ 18MG BASE</u>	<u>A090609 002</u>	Feb 23, 2018
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>A090609 003</u>	Feb 23, 2018
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A090609 004</u>	Feb 23, 2018
<u>AB</u>		<u>EQ 60MG BASE</u>	<u>A090609 005</u>	Feb 23, 2018
<u>AB</u>		<u>EQ 80MG BASE</u>	<u>A090609 006</u>	Feb 23, 2018
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A090609 007</u>	Feb 23, 2018
<u>AB</u>	GLENMARK PHARMS LTD	<u>EQ 10MG BASE</u>	<u>A079019 001</u>	May 30, 2017
<u>AB</u>		<u>EQ 18MG BASE</u>	<u>A079019 002</u>	May 30, 2017
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>A079019 003</u>	May 30, 2017
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A079019 004</u>	May 30, 2017
<u>AB</u>		<u>EQ 60MG BASE</u>	<u>A079019 005</u>	May 30, 2017
<u>AB</u>		<u>EQ 80MG BASE</u>	<u>A079019 006</u>	May 30, 2017
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A079019 007</u>	May 30, 2017
<u>AB</u>	HETERO LABS LTD V	<u>EQ 10MG BASE</u>	<u>A202682 001</u>	Mar 11, 2021
<u>AB</u>		<u>EQ 18MG BASE</u>	<u>A202682 002</u>	Mar 11, 2021
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>A202682 003</u>	Mar 11, 2021
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A202682 004</u>	Mar 11, 2021
<u>AB</u>		<u>EQ 60MG BASE</u>	<u>A202682 005</u>	Mar 11, 2021
<u>AB</u>		<u>EQ 80MG BASE</u>	<u>A202682 006</u>	Mar 11, 2021
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A202682 007</u>	Mar 11, 2021
<u>AB</u>	STRIDES PHARMA	<u>EQ 10MG BASE</u>	<u>A079021 001</u>	Feb 18, 2021
<u>AB</u>		<u>EQ 18MG BASE</u>	<u>A079021 002</u>	Feb 18, 2021
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>A079021 003</u>	Feb 18, 2021
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A079021 004</u>	Feb 18, 2021
<u>AB</u>		<u>EQ 60MG BASE</u>	<u>A079021 005</u>	Feb 18, 2021
<u>AB</u>		<u>EQ 80MG BASE</u>	<u>A079021 006</u>	Feb 18, 2021
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A079021 007</u>	Feb 18, 2021
<u>AB</u>	TEVA PHARMS USA	<u>EQ 10MG BASE</u>	<u>A079022 001</u>	May 30, 2017
<u>AB</u>		<u>EQ 18MG BASE</u>	<u>A079022 002</u>	May 30, 2017
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>A079022 003</u>	May 30, 2017
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A079022 004</u>	May 30, 2017
<u>AB</u>		<u>EQ 60MG BASE</u>	<u>A079022 005</u>	May 30, 2017
<u>AB</u>		<u>EQ 80MG BASE</u>	<u>A079022 006</u>	May 30, 2017
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A079022 007</u>	May 30, 2017
<u>AB</u>	ZYDUS PHARMS USA INC	<u>EQ 10MG BASE</u>	<u>A079017 007</u>	Apr 05, 2023
<u>AB</u>		<u>EQ 18MG BASE</u>	<u>A079017 001</u>	Sep 16, 2010
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>A079017 002</u>	Sep 16, 2010
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A079017 003</u>	Sep 16, 2010
<u>AB</u>		<u>EQ 60MG BASE</u>	<u>A079017 004</u>	Sep 16, 2010
<u>AB</u>		<u>EQ 80MG BASE</u>	<u>A079017 005</u>	Sep 16, 2010
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A079017 006</u>	Sep 16, 2010

STRATTERA

<u>AB</u>	+	LILLY	<u>EQ 10MG BASE</u>	<u>N021411 002</u>	Nov 26, 2002
<u>AB</u>	+		<u>EQ 18MG BASE</u>	<u>N021411 003</u>	Nov 26, 2002
<u>AB</u>	+		<u>EQ 25MG BASE</u>	<u>N021411 004</u>	Nov 26, 2002
<u>AB</u>	+		<u>EQ 40MG BASE</u>	<u>N021411 005</u>	Nov 26, 2002
<u>AB</u>	+	!	<u>EQ 60MG BASE</u>	<u>N021411 006</u>	Nov 26, 2002
<u>AB</u>	+		<u>EQ 80MG BASE</u>	<u>N021411 007</u>	Feb 14, 2005
<u>AB</u>	+		<u>EQ 100MG BASE</u>	<u>N021411 008</u>	Feb 14, 2005

ATORVASTATIN CALCIUM

SUSPENSION; ORAL

ATORVALIQ

#! CMP DEV LLC

20MG/5ML

N213260 001 Feb 01, 2023

TABLET; ORAL

ATORVASTATIN CALCIUM

<u>AB</u>	ACCORD HLTHCARE	<u>EQ 10MG BASE</u>	<u>A207687 001</u>	Mar 30, 2018
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A207687 002</u>	Mar 30, 2018
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A207687 003</u>	Mar 30, 2018
<u>AB</u>		<u>EQ 80MG BASE</u>	<u>A207687 004</u>	Mar 30, 2018
<u>AB</u>	AGNITIO	<u>EQ 10MG BASE</u>	<u>A214969 001</u>	Sep 02, 2021
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A214969 002</u>	Sep 02, 2021
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A214969 003</u>	Sep 02, 2021

PRESCRIPTION DRUG PRODUCT LIST

ATORVASTATIN CALCIUM

TABLET; ORAL

ATORVASTATIN CALCIUM

<u>AB</u>		<u>EQ 80MG BASE</u>	<u>A214969 004</u>	Sep 02, 2021
<u>AB</u>	ALKEM LABS LTD	<u>EQ 10MG BASE</u>	<u>A209288 001</u>	Dec 21, 2018
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A209288 002</u>	Dec 21, 2018
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A209288 003</u>	Dec 21, 2018
<u>AB</u>		<u>EQ 80MG BASE</u>	<u>A209288 004</u>	Dec 21, 2018
<u>AB</u>	APOTEX INC	<u>EQ 10MG BASE</u>	<u>A090548 001</u>	May 29, 2012
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A090548 002</u>	May 29, 2012
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A090548 003</u>	May 29, 2012
<u>AB</u>		<u>EQ 80MG BASE</u>	<u>A090548 004</u>	May 29, 2012
<u>AB</u>	AUROBINDO PHARMA LTD	<u>EQ 10MG BASE</u>	<u>A217118 001</u>	Feb 13, 2024
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A217118 002</u>	Feb 13, 2024
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A217118 003</u>	Feb 13, 2024
<u>AB</u>		<u>EQ 80MG BASE</u>	<u>A217118 004</u>	Feb 13, 2024
<u>AB</u>	BIOCON PHARMA	<u>EQ 10MG BASE</u>	<u>A216436 001</u>	Nov 23, 2022
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A216436 002</u>	Nov 23, 2022
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A216436 003</u>	Nov 23, 2022
<u>AB</u>		<u>EQ 80MG BASE</u>	<u>A216436 004</u>	Nov 23, 2022
<u>AB</u>	CHARTWELL RX	<u>EQ 10MG BASE</u>	<u>A077575 001</u>	May 29, 2012
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A077575 002</u>	May 29, 2012
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A077575 003</u>	May 29, 2012
<u>AB</u>		<u>EQ 80MG BASE</u>	<u>A077575 004</u>	May 29, 2012
<u>AB</u>	DR REDDYS	<u>EQ 10MG BASE</u>	<u>A214659 001</u>	Jul 14, 2021
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A214659 002</u>	Jul 14, 2021
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A214659 003</u>	Jul 14, 2021
<u>AB</u>		<u>EQ 80MG BASE</u>	<u>A214659 004</u>	Jul 14, 2021
<u>AB</u>	DR REDDYS LABS LTD	<u>EQ 10MG BASE</u>	<u>A091650 001</u>	Jul 17, 2012
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A091650 002</u>	Jul 17, 2012
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A091650 003</u>	Jul 17, 2012
<u>AB</u>		<u>EQ 80MG BASE</u>	<u>A202357 001</u>	Jul 17, 2012
<u>AB</u>	GRAVITI PHARMS	<u>EQ 10MG BASE</u>	<u>A209912 001</u>	Jun 18, 2018
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A209912 002</u>	Jun 18, 2018
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A209912 003</u>	Jun 18, 2018
<u>AB</u>		<u>EQ 80MG BASE</u>	<u>A209912 004</u>	Jun 18, 2018
<u>AB</u>	HETERO LABS LTD V	<u>EQ 10MG BASE</u>	<u>A214344 001</u>	Sep 12, 2023
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A214344 002</u>	Sep 12, 2023
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A214344 003</u>	Sep 12, 2023
<u>AB</u>		<u>EQ 80MG BASE</u>	<u>A214344 004</u>	Sep 12, 2023
<u>AB</u>	LANNETT CO INC	<u>EQ 10MG BASE</u>	<u>A091624 001</u>	Apr 05, 2013
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A091624 002</u>	Apr 05, 2013
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A091624 003</u>	Apr 05, 2013
<u>AB</u>		<u>EQ 80MG BASE</u>	<u>A091624 004</u>	Apr 05, 2013
<u>AB</u>	LAURUS	<u>EQ 10MG BASE</u>	<u>A214513 001</u>	Jan 22, 2024
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A214513 002</u>	Jan 22, 2024
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A214513 003</u>	Jan 22, 2024
<u>AB</u>		<u>EQ 80MG BASE</u>	<u>A214513 004</u>	Jan 22, 2024
<u>AB</u>	MACLEODS PHARMS LTD	<u>EQ 10MG BASE</u>	<u>A208480 001</u>	Jun 03, 2024
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A208480 002</u>	Jun 03, 2024
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A208480 003</u>	Jun 03, 2024
<u>AB</u>		<u>EQ 80MG BASE</u>	<u>A208480 004</u>	Jun 03, 2024
<u>AB</u>	MANKIND PHARMA	<u>EQ 10MG BASE</u>	<u>A217081 001</u>	Jan 05, 2024
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A217081 002</u>	Jan 05, 2024
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A217081 003</u>	Jan 05, 2024
<u>AB</u>		<u>EQ 80MG BASE</u>	<u>A217081 004</u>	Jan 05, 2024
<u>AB</u>	MICRO LABS LTD INDIA	<u>EQ 10MG BASE</u>	<u>A205945 001</u>	Nov 07, 2019
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A205945 002</u>	Nov 07, 2019
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A205945 003</u>	Nov 07, 2019
<u>AB</u>		<u>EQ 80MG BASE</u>	<u>A205945 004</u>	Nov 07, 2019
<u>AB</u>	MSN	<u>EQ 10MG BASE</u>	<u>A211933 001</u>	Feb 08, 2019
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A211933 002</u>	Feb 08, 2019
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A211933 003</u>	Feb 08, 2019
<u>AB</u>		<u>EQ 80MG BASE</u>	<u>A211933 004</u>	Feb 08, 2019
<u>AB</u>	MYLAN PHARMS INC	<u>EQ 10MG BASE</u>	<u>A091226 001</u>	May 29, 2012
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A091226 002</u>	May 29, 2012
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A091226 003</u>	May 29, 2012
<u>AB</u>		<u>EQ 80MG BASE</u>	<u>A091226 004</u>	May 29, 2012
<u>AB</u>	SCIEGEN PHARMS INC	<u>EQ 10MG BASE</u>	<u>A205519 001</u>	May 19, 2016
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A205519 002</u>	May 19, 2016
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A205519 003</u>	May 19, 2016

PRESCRIPTION DRUG PRODUCT LIST

ATORVASTATIN CALCIUM

TABLET; ORAL

ATORVASTATIN CALCIUM

<u>AB</u>		<u>EQ 80MG BASE</u>	<u>A205519 004</u>	May 19, 2016
<u>AB</u>	SHANDONG XINHUA	<u>EQ 10MG BASE</u>	<u>A211886 001</u>	Jan 25, 2024
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A211886 002</u>	Jan 25, 2024
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A211886 003</u>	Jan 25, 2024
<u>AB</u>		<u>EQ 80MG BASE</u>	<u>A211886 004</u>	Jan 25, 2024
<u>AB</u>	SUN PHARM INDS LTD	<u>EQ 10MG BASE</u>	<u>A076477 001</u>	Nov 30, 2011
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A076477 002</u>	Nov 30, 2011
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A076477 003</u>	Nov 30, 2011
<u>AB</u>		<u>EQ 80MG BASE</u>	<u>A076477 004</u>	Nov 30, 2011
<u>AB</u>	TEVA PHARMS USA	<u>EQ 10MG BASE</u>	<u>A205300 001</u>	Mar 27, 2017
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A205300 002</u>	Mar 27, 2017
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A205300 003</u>	Mar 27, 2017
<u>AB</u>		<u>EQ 80MG BASE</u>	<u>A205300 004</u>	Mar 27, 2017
<u>AB</u>	ZYDUS PHARMS	<u>EQ 10MG BASE</u>	<u>A206536 001</u>	Nov 20, 2018
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A206536 002</u>	Nov 20, 2018
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A206536 003</u>	Nov 20, 2018
<u>AB</u>		<u>EQ 80MG BASE</u>	<u>A206536 004</u>	Nov 20, 2018

LIPITOR

<u>AB</u>	+ UPJOHN	<u>EQ 10MG BASE</u>	<u>N020702 001</u>	Dec 17, 1996
<u>AB</u>	+	<u>EQ 20MG BASE</u>	<u>N020702 002</u>	Dec 17, 1996
<u>AB</u>	+	<u>EQ 40MG BASE</u>	<u>N020702 003</u>	Dec 17, 1996
<u>AB</u>	+!	<u>EQ 80MG BASE</u>	<u>N020702 004</u>	Apr 07, 2000

ATORVASTATIN CALCIUM

BX	UMEDICA	EQ 10MG BASE	A213853 001	Aug 19, 2020
BX		EQ 20MG BASE	A213853 002	Aug 19, 2020
BX		EQ 40MG BASE	A213853 003	Aug 19, 2020
BX		EQ 80MG BASE	A213853 004	Aug 19, 2020

ATORVASTATIN CALCIUM; EZETIMIBE

TABLET; ORAL

LYPQOZET

	ALThERA PHARMS	EQ 10MG BASE;10MG	A206084 001	Apr 26, 2017
		EQ 20MG BASE;10MG	A206084 002	Apr 26, 2017
		EQ 40MG BASE;10MG	A206084 003	Apr 26, 2017
	!	EQ 80MG BASE;10MG	A206084 004	Apr 26, 2017

ATOVAQUONE

SUSPENSION; ORAL

ATOVAQUONE

<u>AB</u>	ABHAI LLC	<u>750MG/5ML</u>	<u>A210510 001</u>	May 31, 2019
<u>AB</u>	ABON PHARMS LLC	<u>750MG/5ML</u>	<u>A214272 001</u>	Oct 25, 2021
<u>AB</u>	AMNEAL PHARMS	<u>750MG/5ML</u>	<u>A202960 001</u>	Mar 18, 2014
<u>AB</u>	APOTEX	<u>750MG/5ML</u>	<u>A209750 001</u>	Oct 11, 2017
<u>AB</u>	BIONPHARMA	<u>750MG/5ML</u>	<u>A212918 001</u>	Mar 30, 2021
<u>AB</u>	CHARTWELL RX	<u>750MG/5ML</u>	<u>A207833 001</u>	Apr 28, 2017
<u>AB</u>	GLENMARK SPECLT	<u>750MG/5ML</u>	<u>A209685 001</u>	Nov 21, 2018
<u>AB</u>	HETERO LABS LTD III	<u>750MG/5ML</u>	<u>A210692 001</u>	Oct 11, 2018
<u>AB</u>	LUPIN LTD	<u>750MG/5ML</u>	<u>A209105 001</u>	Sep 11, 2018

MEPRON

<u>AB</u>	+!	GLAXOSMITHKLINE LLC	<u>750MG/5ML</u>	<u>N020500 001</u>	Feb 08, 1995
-----------	----	---------------------	------------------	--------------------	--------------

ATOVAQUONE; PROGUANIL HYDROCHLORIDE

TABLET; ORAL

ATOVAQUONE AND PROGUANIL HYDROCHLORIDE

<u>AB</u>	GLENMARK PHARMS LTD	<u>62.5MG;25MG</u>	<u>A091211 002</u>	Apr 06, 2015
<u>AB</u>		<u>250MG;100MG</u>	<u>A091211 001</u>	Jan 12, 2011
<u>AB</u>	MYLAN	<u>62.5MG;25MG</u>	<u>A202362 001</u>	May 27, 2014
<u>AB</u>		<u>250MG;100MG</u>	<u>A202362 002</u>	May 27, 2014

MALARONE

<u>AB</u>	+!	GLAXOSMITHKLINE	<u>250MG;100MG</u>	<u>N021078 001</u>	Jul 14, 2000
-----------	----	-----------------	--------------------	--------------------	--------------

MALARONE PEDIATRIC

<u>AB</u>	+	GLAXOSMITHKLINE	<u>62.5MG;25MG</u>	<u>N021078 002</u>	Jul 14, 2000
-----------	---	-----------------	--------------------	--------------------	--------------

ATRACURIUM BESYLATE

INJECTABLE; INJECTION

ATRACURIUM BESYLATE

<u>AP</u>	EUGIA PHARMA	<u>10MG/ML</u>	<u>A206011 001</u>	Apr 08, 2015
<u>AP</u>	!	HIKMA	<u>A074901 001</u>	Jul 18, 1997
<u>AP</u>	HOSPIRA INC	<u>10MG/ML</u>	<u>A090761 001</u>	Oct 18, 2012
<u>AP</u>	MEITHEAL	<u>10MG/ML</u>	<u>A091489 001</u>	Feb 17, 2012

PRESCRIPTION DRUG PRODUCT LIST

ATRACURIUM BESYLATE

INJECTABLE; INJECTION

ATRACURIUM BESYLATE PRESERVATIVE FREE

<u>AP</u>	EUGIA PHARMA	<u>10MG/ML</u>	<u>A206010 001</u>	Apr 08, 2015
<u>AP</u>	! HIKMA	<u>10MG/ML</u>	<u>A074900 001</u>	Jul 18, 1997
<u>AP</u>	HOSPIRA INC	<u>10MG/ML</u>	<u>A090782 001</u>	Oct 18, 2012
<u>AP</u>	MEITHEAL	<u>10MG/ML</u>	<u>A091488 001</u>	Feb 17, 2012

ATROPINE SULFATE

SOLUTION; INTRAVENOUS

ATROPINE SULFATE

<u>AP</u>	ACCORD HLTHCARE	<u>0.25MG/5ML (0.05MG/ML)</u>	<u>A212868 001</u>	Jul 26, 2021
<u>AP</u>	+!	<u>0.4MG/ML (0.4MG/ML)</u>	<u>N214652 001</u>	Sep 29, 2020
<u>AP</u>		<u>0.5MG/5ML (0.1MG/ML)</u>	<u>A212868 002</u>	Jul 26, 2021
<u>AP</u>		<u>1MG/10ML (0.1MG/ML)</u>	<u>A212868 003</u>	Jul 26, 2021
<u>AP</u>	+!	<u>1MG/ML (1MG/ML)</u>	<u>N214652 002</u>	Sep 29, 2020
<u>AP</u>	AM REGENT	<u>0.4MG/ML (0.4MG/ML)</u>	<u>A216120 001</u>	May 26, 2022
<u>AP</u>		<u>1MG/ML (1MG/ML)</u>	<u>A216120 002</u>	May 26, 2022
<u>AP</u>	AMNEAL	<u>0.5MG/5ML (0.1MG/ML)</u>	<u>A215342 001</u>	Jan 26, 2022
<u>AP</u>		<u>1MG/10ML (0.1MG/ML)</u>	<u>A215342 002</u>	Oct 25, 2023
<u>AP</u>	+! HOSPIRA	<u>0.25MG/5ML (0.05MG/ML)</u>	<u>N021146 002</u>	Jul 09, 2001
<u>AP</u>	+!	<u>0.5MG/5ML (0.1MG/ML)</u>	<u>N021146 001</u>	Jul 09, 2001
<u>AP</u>	+!	<u>1MG/10ML (0.1MG/ML)</u>	<u>N021146 003</u>	Jul 09, 2001
<u>AP</u>	INTL MEDICATION SYS	<u>1MG/10ML (0.1MG/ML)</u>	<u>A212461 001</u>	Oct 05, 2020
<u>AP</u>	MEDEFIL INC	<u>1MG/10ML (0.1MG/ML)</u>	<u>A214970 001</u>	Nov 04, 2022
<u>AP</u>	SOMERSET THERAPS LLC	<u>0.4MG/ML (0.4MG/ML)</u>	<u>A215969 001</u>	Jul 03, 2024
<u>AP</u>		<u>1MG/ML (1MG/ML)</u>	<u>A215969 002</u>	Jul 03, 2024

SOLUTION; INTRAVENOUS, INTRAMUSCULAR, SUBCUTANEOUS, INTRAOSSEOUS, ENDOTRACHEAL

ATROPINE SULFATE

<u>AP</u>	ACCORD HLTHCARE	<u>8MG/20ML (0.4MG/ML)</u>	<u>A213424 001</u>	Mar 19, 2021
<u>AP</u>	+! FRESENIUS KABI USA	<u>8MG/20ML (0.4MG/ML)</u>	<u>N209260 001</u>	Jan 26, 2018
<u>AP</u>	HIKMA	<u>8MG/20ML (0.4MG/ML)</u>	<u>A213561 001</u>	Dec 01, 2021
<u>AP</u>	SOMERSET THERAPS LLC	<u>8MG/20ML (0.4MG/ML)</u>	<u>A215005 001</u>	Nov 29, 2024

SOLUTION/DROPS; OPHTHALMIC

ATROPINE SULFATE

<u>AT1</u>	! AMNEAL	<u>1%</u>	<u>A214752 001</u>	Jul 14, 2022
<u>AT1</u>	APOTEX	<u>1%</u>	<u>A215624 001</u>	Nov 26, 2021
<u>AT1</u>	MANKIND PHARMA	<u>1%</u>	<u>A218148 001</u>	Jan 08, 2024
<u>AT1</u>	+ RISING	<u>1%</u>	<u>N206289 001</u>	Jul 18, 2014
<u>AT1</u>	SOMERSET THERAPS LLC	<u>1%</u>	<u>A215618 001</u>	Jul 01, 2024
<u>AT2</u>	SOMERSET	<u>1%</u>	<u>A217791 001</u>	Apr 29, 2024

ISOPTO ATROPINE

<u>AT2</u>	+! ALCON LABS INC	<u>1%</u>	<u>N208151 001</u>	Dec 01, 2016
------------	-------------------	-----------	--------------------	--------------

ATROPINE SULFATE

	+! BAUSCH AND LOMB INC	<u>1%</u>	<u>N213581 001</u>	Mar 15, 2022
--	------------------------	-----------	--------------------	--------------

ATROPINE SULFATE; DIFENOXIN HYDROCHLORIDE

TABLET; ORAL

MOTOFEN

	+! LEGACY PHARMA	<u>0.025MG; 1MG</u>	<u>N017744 002</u>	
--	------------------	---------------------	--------------------	--

ATROPINE SULFATE; DIPHENOXYLATE HYDROCHLORIDE

SOLUTION; ORAL

DIPHENOXYLATE HYDROCHLORIDE AND ATROPINE SULFATE

!	HIKMA	<u>0.025MG/5ML; 2.5MG/5ML</u>	<u>A087708 001</u>	May 03, 1982
---	-------	-------------------------------	--------------------	--------------

TABLET; ORAL

DIPHENOXYLATE HYDROCHLORIDE AND ATROPINE SULFATE

<u>AA</u>	CHARTWELL RX	<u>0.025MG; 2.5MG</u>	<u>A207128 001</u>	Oct 21, 2020
<u>AA</u>	LANNETT	<u>0.025MG; 2.5MG</u>	<u>A085372 001</u>	
<u>AA</u>	LEADING	<u>0.025MG; 2.5MG</u>	<u>A213413 001</u>	Feb 20, 2020
<u>AA</u>	SPECGX LLC	<u>0.025MG; 2.5MG</u>	<u>A213335 001</u>	Oct 06, 2020
<u>AA</u>	UNICHEM	<u>0.025MG; 2.5MG</u>	<u>A210819 001</u>	Nov 13, 2018
<u>AA</u>	WINDER LABS LLC	<u>0.025MG; 2.5MG</u>	<u>A211362 001</u>	Jan 27, 2021

LOMOTIL

<u>AA</u>	+! PFIZER	<u>0.025MG; 2.5MG</u>	<u>N012462 001</u>	
-----------	-----------	-----------------------	--------------------	--

PRESCRIPTION DRUG PRODUCT LISTATROPINE; PRALIDOXIME CHLORIDE

INJECTABLE; INTRAMUSCULAR

DUODOTE

+! MMT 2.1MG/0.7ML; 600MG/2ML N021983 001 Sep 28, 2006

AURANOFIN

CAPSULE; ORAL

RIDAURA

+! LEGACY PHARMA 3MG N018689 001 May 24, 1985

AVACINCAPTAD PEGOL SODIUM

SOLUTION; INTRAVITREAL

IZERVAY

+! ASTELLAS EQ 2MG BASE/0.1ML (EQ 2MG BASE/0.1ML) N217225 001 Aug 04, 2023

AVACOPAN

CAPSULE; ORAL

TAVNEOS

+! CHEMOCENTRYX 10MG N214487 001 Oct 07, 2021

AVANAFIL

TABLET; ORAL

AVANAFIL**AB** HETERO LABS LTD V **50MG** **A209266 001** Jun 14, 2024**AB** **100MG** **A209266 002** Jun 14, 2024**AB** **200MG** **A209266 003** Jun 14, 2024STENDRA**AB** + METUCHEN PHARMS **50MG** **N202276 001** Apr 27, 2012**AB** + **100MG** **N202276 002** Apr 27, 2012**AB** +! **200MG** **N202276 003** Apr 27, 2012AVAPRITINIB

TABLET; ORAL

AYVAKIT

+ BLUEPRINT MEDICINES 25MG N212608 004 Jun 16, 2021

+ 50MG N212608 005 Jun 16, 2021

+ 100MG N212608 001 Jan 09, 2020

+ 200MG N212608 002 Jan 09, 2020

+! 300MG N212608 003 Jan 09, 2020

AVATROMBOPAG MALEATE

TABLET; ORAL

DOPTELET

+! AKARX INC EQ 20MG BASE N210238 001 May 21, 2018

AVIBACTAM SODIUM; CEFTAZIDIME

POWDER; INTRAVENOUS

AVYCAZ

+! ABBVIE EQ 0.5GM BASE; 2GM/VIAL N206494 001 Feb 25, 2015

AXITINIB

TABLET; ORAL

INLYTA

+ PF PRISM CV 1MG N202324 001 Jan 27, 2012

+! 5MG N202324 002 Jan 27, 2012

AZACITIDINE

POWDER; INTRAVENOUS, SUBCUTANEOUS

AZACITIDINE**AP** ACCORD HLTHCARE **100MG/VIAL** **A207475 001** Jul 02, 2018**AP** ACTAVIS LLC **100MG/VIAL** **N208216 001** Apr 29, 2016**AP** AMNEAL **100MG/VIAL** **A211549 001** Feb 03, 2022**AP** CIPLA **100MG/VIAL** **A209540 001** May 04, 2018**AP** DR REDDYS **100MG/VIAL** **A201537 001** Sep 16, 2013**AP** EUGIA PHARMA **100MG/VIAL** **A215066 001** Dec 30, 2022**AP** EUROHLTH INTL SARL **100MG/VIAL** **A209337 001** Jun 08, 2020**AP** HETERO LABS LTD VI **100MG/VIAL** **A215765 001** Oct 15, 2024**AP** JIANGSU HANSOH **100MG/VIAL** **A215905 001** Jun 28, 2023

PHARM

AP MEITHEAL **100MG/VIAL** **A212128 001** Nov 02, 2020**AP** MSN LABS PVT LTD **100MG/VIAL** **A212580 001** May 16, 2024**AP** NATCO PHARMA LTD **100MG/VIAL** **A207234 001** Jun 23, 2017**AP** SHILPA MEDICARE **100MG/VIAL** **A207518 001** Sep 29, 2016VIDAZA**AP** +! BRISTOL-MYERS **100MG/VIAL** **N050794 001** May 19, 2004

PRESCRIPTION DRUG PRODUCT LIST

AZACITIDINE

TABLET; ORAL

ONUREG

+ BRISTOL

200MG

N214120 001 Sep 01, 2020

+!

300MG

N214120 002 Sep 01, 2020

AZATHIOPRINE

TABLET; ORAL

AZASAN**AB** AAIPHARMA LLC**25MG****A075252 002** Feb 03, 2003**AB****50MG****A075252 001** Jun 07, 1999**AB****75MG****A075252 003** Feb 03, 2003**AB****100MG****A075252 004** Feb 03, 2003AZATHIOPRINE**AB** ALKEM LABS LTD**25MG****A208687 001** Mar 27, 2020**AB****50MG****A208687 002** Mar 27, 2020**AB****75MG****A208687 003** Mar 27, 2020**AB****100MG****A208687 004** Mar 27, 2020**AB** AMNEAL**50MG****A074069 001** Feb 16, 1996**AB****75MG****A074069 002** Nov 02, 2021**AB****100MG****A074069 003** Nov 02, 2021**AB** RISING**50MG****A075568 001** Dec 13, 1999**AB**

ZYDUS PHARMS USA

25MG**A077621 002** Sep 05, 2008**AB****50MG****A077621 001** Mar 15, 2007**AB****75MG****A077621 003** Sep 05, 2008**AB****100MG****A077621 004** Sep 05, 2008TMURAN**AB** +! LEGACY PHARMA**50MG****N016324 001**AZATHIOPRINE SODIUM

INJECTABLE; INJECTION

AZATHIOPRINE SODIUM

! HIKMA

EQ 100MG BASE/VIAL

A074419 001 Mar 31, 1995

AZELAIC ACID

AEROSOL, FOAM; TOPICAL

FINACEA

+! LEO PHARMA AS

15%

N207071 001 Jul 29, 2015

CREAM; TOPICAL

AZELEX

+! ALMIRALL

20%

N020428 001 Sep 13, 1995

GEL; TOPICAL

AZELAIC ACID**AB** ACTAVIS LABS UT INC**15%****A208011 001** Nov 19, 2018**AB**

ENCUBE

15%**A208724 001** Nov 19, 2018**AB**

GLENMARK SPECLT

15%**A204637 001** Nov 19, 2018**AB**

TARO

15%**A210549 001** Aug 23, 2019FINACEA**AB** +! LEO PHARMA AS**15%****N021470 001** Dec 24, 2002AZELASTINE HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

AZELASTINE HYDROCHLORIDE**AT** ALEMBIC**0.05%****A209620 001** Mar 20, 2019**AT**

APOTEX INC

0.05%**A078621 001** Aug 03, 2009**AT**

GLAND PHARMA LTD

0.05%**A210092 001** Feb 25, 2020**AT**

! SANDOZ

0.05%**A202305 001** May 31, 2012**AT**

SOMERSET THERAPS

0.05%**A207411 001** Mar 29, 2019

LLC

SPRAY, METERED; NASAL

AZELASTINE HYDROCHLORIDE**AB** ALKEM LABS LTD**0.137MG/SPRAY****A208156 001** Aug 18, 2017**AB**

AMNEAL

0.137MG/SPRAY**A204660 001** Aug 28, 2017**AB**

! APOTEX INC

0.137MG/SPRAY**A077954 001** Apr 30, 2009**AB**

AUROBINDO PHARMA

0.137MG/SPRAY**A212289 001** May 08, 2020

LTD

AB BIONPHARMA**0.137MG/SPRAY****A090176 001** Jul 28, 2015**AB**

EPIC PHARMA LLC

0.137MG/SPRAY**A207610 001** May 17, 2019**AB**

UPSHER SMITH LABS

0.137MG/SPRAY**A202609 001** Mar 17, 2017**AB**

ZYDUS PHARMS

0.137MG/SPRAY**A091409 001** Aug 14, 2017

PRESCRIPTION DRUG PRODUCT LIST

AZELASTINE HYDROCHLORIDE; FLUTICASONE PROPIONATE

SPRAY, METERED;NASAL

AZELASTINE HYDROCHLORIDE AND FLUTICASONE PROPIONATE

AB	!	APOTEX	<u>0.137MG/SPRAY;0.05MG/SPRAY</u>	<u>A207712 001</u>	Apr 28, 2017
AB		PADAGIS ISRAEL	<u>0.137MG/SPRAY;0.05MG/SPRAY</u>	<u>A208111 001</u>	Feb 18, 2021

DYMISTA

AB	+	MYLAN SPECIALITY LP	<u>0.137MG/SPRAY;0.05MG/SPRAY</u>	<u>N202236 001</u>	May 01, 2012
-----------	----------	---------------------	--	---------------------------	--------------

AZILSARTAN KAMEDOXOMIL

TABLET;ORAL

EDARBI

+	AZURITY	EQ 40MG MEDOXOMIL	N200796 001	Feb 25, 2011
+	!	EQ 80MG MEDOXOMIL	N200796 002	Feb 25, 2011

AZILSARTAN KAMEDOXOMIL; CHLORTHALIDONE

TABLET;ORAL

EDARBYCLOR

+	AZURITY	EQ 40MG MEDOXOMIL;12.5MG	N202331 001	Dec 20, 2011
+	!	EQ 40MG MEDOXOMIL;25MG	N202331 002	Dec 20, 2011

AZITHROMYCIN

FOR SUSPENSION;ORAL

AZITHROMYCIN

AB		AMNEAL	<u>EQ 100MG BASE/5ML</u>	<u>A205666 001</u>	Jul 19, 2018
AB			<u>EQ 200MG BASE/5ML</u>	<u>A205666 002</u>	Jul 19, 2018
AB		AUROBINDO PHARMA LTD	<u>EQ 100MG BASE/5ML</u>	<u>A209201 001</u>	Oct 09, 2018
AB			<u>EQ 200MG BASE/5ML</u>	<u>A209201 002</u>	Oct 09, 2018
AB		EPIC PHARMA LLC	<u>EQ 100MG BASE/5ML</u>	<u>A207531 001</u>	Apr 09, 2018
AB			<u>EQ 200MG BASE/5ML</u>	<u>A207531 002</u>	Apr 09, 2018
AB		HAINAN POLY	<u>EQ 100MG BASE/5ML</u>	<u>A217036 001</u>	Jul 27, 2023
AB			<u>EQ 200MG BASE/5ML</u>	<u>A217036 002</u>	Jul 27, 2023
AB		PLIVA	<u>EQ 100MG BASE/5ML</u>	<u>A065246 002</u>	Jul 05, 2006
AB			<u>EQ 200MG BASE/5ML</u>	<u>A065246 001</u>	Jul 05, 2006
AB		ZYDUS LIFESCIENCES	<u>EQ 100MG BASE/5ML</u>	<u>A211147 001</u>	Jul 31, 2018
AB			<u>EQ 200MG BASE/5ML</u>	<u>A211147 002</u>	Jul 31, 2018

ZITHROMAX

AB	+	PFIZER	<u>EQ 100MG BASE/5ML</u>	<u>N050710 001</u>	Oct 19, 1995
AB	+	!	<u>EQ 200MG BASE/5ML</u>	<u>N050710 002</u>	Oct 19, 1995
		!	EQ 1GM BASE/PACKET	N050693 001	Sep 28, 1994

INJECTABLE;INJECTION

AZITHROMYCIN

AP		EUGIA PHARMA	<u>EQ 500MG BASE/VIAL</u>	<u>A203294 001</u>	Jun 19, 2015
AP		FRESENIUS KABI USA	<u>EQ 500MG BASE/VIAL</u>	<u>A065179 001</u>	Dec 13, 2005
AP		GLAND PHARMA LTD	<u>EQ 500MG BASE/VIAL</u>	<u>A065501 001</u>	Nov 09, 2009
AP		HOSPIRA	<u>EQ 500MG BASE/VIAL</u>	<u>A065500 001</u>	Jun 26, 2009
AP			<u>EQ 500MG BASE/VIAL</u>	<u>A065511 001</u>	Jun 26, 2009
AP		SLATE RUN PHARMA	<u>EQ 500MG BASE/VIAL</u>	<u>A203412 001</u>	Oct 09, 2018

ZITHROMAX

AP	+	PFIZER	<u>EQ 500MG BASE/VIAL</u>	<u>N050733 001</u>	Jan 30, 1997
-----------	----------	--------	----------------------------------	---------------------------	--------------

SOLUTION/DROPS;OPHTHALMIC

+	!	THEA PHARMA	1%	N050810 001	Apr 27, 2007
----------	---	-------------	----	-------------	--------------

TABLET;ORAL

AZITHROMYCIN

AB		ALEMBIC	<u>EQ 250MG BASE</u>	<u>A211791 001</u>	Jan 28, 2020
AB			<u>EQ 500MG BASE</u>	<u>A211792 001</u>	Jan 28, 2020
AB			<u>EQ 600MG BASE</u>	<u>A211793 001</u>	Jan 27, 2020
AB		AUROBINDO PHARMA LTD	<u>EQ 250MG BASE</u>	<u>A207370 001</u>	Jul 05, 2018
AB			<u>EQ 500MG BASE</u>	<u>A207398 001</u>	Jul 05, 2018
AB		BIONPHARMA	<u>EQ 250MG BASE</u>	<u>A210000 001</u>	Feb 26, 2019
AB			<u>EQ 500MG BASE</u>	<u>A210001 001</u>	Feb 26, 2019
AB			<u>EQ 600MG BASE</u>	<u>A209999 001</u>	Dec 26, 2018
AB		CHARTWELL RX	<u>EQ 250MG BASE</u>	<u>A065404 001</u>	Feb 11, 2008
AB			<u>EQ 500MG BASE</u>	<u>A065405 001</u>	Feb 11, 2008
AB			<u>EQ 600MG BASE</u>	<u>A065302 003</u>	Feb 11, 2008
AB		CSPC OUYI	<u>EQ 250MG BASE</u>	<u>A208250 001</u>	Apr 17, 2019
AB			<u>EQ 500MG BASE</u>	<u>A208249 001</u>	Oct 25, 2018
AB			<u>EQ 600MG BASE</u>	<u>A207566 001</u>	Sep 24, 2018
AB		LUPIN LTD	<u>EQ 250MG BASE</u>	<u>A065398 001</u>	May 15, 2015
AB			<u>EQ 500MG BASE</u>	<u>A065399 001</u>	May 15, 2015
AB			<u>EQ 600MG BASE</u>	<u>A065400 001</u>	May 15, 2015
AB		PLIVA	<u>EQ 250MG BASE</u>	<u>A065225 001</u>	Nov 14, 2005

PRESCRIPTION DRUG PRODUCT LIST

AZITHROMYCIN

TABLET; ORAL

AZITHROMYCIN

<u>AB</u>		<u>EQ 500MG BASE</u>	<u>A065223 001</u>	Nov 14, 2005
<u>AB</u>	!	<u>EQ 600MG BASE</u>	<u>A065218 001</u>	Nov 14, 2005
<u>AB</u>	SANDOZ	<u>EQ 250MG BASE</u>	<u>A065211 001</u>	Nov 14, 2005
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>A065212 001</u>	Nov 14, 2005
<u>AB</u>		<u>EQ 600MG BASE</u>	<u>A065209 001</u>	Nov 14, 2005
<u>AB</u>	STRIDES PHARMA	<u>EQ 250MG BASE</u>	<u>A215772 001</u>	Jul 15, 2022
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>A215773 001</u>	Jul 12, 2022
<u>AB</u>	SUNSHINE	<u>EQ 250MG BASE</u>	<u>A209045 001</u>	Dec 07, 2018
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>A209044 001</u>	Dec 07, 2018
<u>AB</u>		<u>EQ 600MG BASE</u>	<u>A209043 001</u>	Dec 06, 2018
<u>AB</u>	TEVA	<u>EQ 500MG BASE</u>	<u>A065193 001</u>	Nov 14, 2005
<u>AB</u>	YUNG SHIN PHARM	<u>EQ 600MG BASE</u>	<u>A211068 001</u>	May 08, 2020

ZITHROMAX

<u>AB</u>	+	PFIZER	<u>EQ 250MG BASE</u>	<u>N050711 001</u>	Jul 18, 1996
<u>AB</u>	+		<u>EQ 500MG BASE</u>	<u>N050784 001</u>	May 24, 2002

AZTREONAM

FOR SOLUTION; INHALATION

CAYSTON

+! GILEAD

75MG/VIAL

N050814 001 Feb 22, 2010

INJECTABLE; INJECTION

AZACTAM

<u>AP</u>	+	BRISTOL MYERS SQUIBB	<u>1GM/VIAL</u>	<u>N050580 002</u>	Dec 31, 1986
<u>AP</u>	+		<u>2GM/VIAL</u>	<u>N050580 003</u>	Dec 31, 1986

AZTREONAM

<u>AP</u>		FRESENIUS KABI USA	<u>1GM/VIAL</u>	<u>A065439 002</u>	Jun 18, 2010
<u>AP</u>			<u>2GM/VIAL</u>	<u>A065439 003</u>	Jun 18, 2010
<u>AP</u>		HOSPIRA	<u>1GM/VIAL</u>	<u>A206517 001</u>	Nov 08, 2021
<u>AP</u>			<u>2GM/VIAL</u>	<u>A206517 002</u>	Nov 08, 2021
		FRESENIUS KABI USA	500MG/VIAL	A065439 001	Jun 18, 2010

BACITRACIN

OINTMENT; OPHTHALMIC

BACITRACIN

! PADAGIS US

500 UNITS/GM

A061212 001

BACITRACIN ZINC; HYDROCORTISONE ACETATE; NEOMYCIN SULFATE; POLYMYXIN B SULFATE

OINTMENT; OPHTHALMIC

BACITRACIN-NEOMYCIN-POLYMYXIN W/ HYDROCORTISONE ACETATE

! PADAGIS US

400 UNITS/GM;1%;EQ 3.5MG BASE/GM;10,000
UNITS/GM

A062166 002

BACITRACIN ZINC; HYDROCORTISONE; NEOMYCIN SULFATE; POLYMYXIN B SULFATE

OINTMENT; OPHTHALMIC

NEOMYCIN AND POLYMYXIN B SULFATES, BACITRACIN ZINC AND HYDROCORTISONE

! BAUSCH AND LOMB

400 UNITS/GM;1%;EQ 3.5MG BASE/GM;10,000
UNITS/GM

A064068 001 Oct 30, 1995

BACITRACIN ZINC; NEOMYCIN SULFATE; POLYMYXIN B SULFATE

OINTMENT; OPHTHALMIC

LUMI-SPORYN

<u>AT</u>	+	CASPER PHARMA LLC	<u>EQ 400 UNITS/GM;EQ 3.5MG BASE/GM;EQ 10,000 UNITS/GM</u>	<u>N050417 001</u>	
			<u>NEOMYCIN AND POLYMYXIN B SULFATES AND BACITRACIN ZINC</u>		
<u>AT</u>	!	BAUSCH AND LOMB	<u>400 UNITS/GM;EQ 3.5MG BASE/GM;10,000 UNITS/GM</u>	<u>A064064 001</u>	Oct 30, 1995
<u>AT</u>		PADAGIS US	<u>400 UNITS/GM;EQ 3.5MG BASE/GM;10,000 UNITS/GM</u>	<u>A060764 002</u>	

BACITRACIN ZINC; POLYMYXIN B SULFATE

OINTMENT; OPHTHALMIC

BACITRACIN ZINC AND POLYMYXIN B SULFATE

<u>AT</u>	!	BAUSCH AND LOMB	<u>500 UNITS/GM;10,000 UNITS/GM</u>	<u>A064046 001</u>	Jan 26, 1995
<u>AT</u>		PADAGIS US	<u>500 UNITS/GM;10,000 UNITS/GM</u>	<u>A065022 001</u>	Feb 27, 2002

BACLOFEN

GRANULES; ORAL

LYVISPAH

+ AMNEAL

5MG/PACKET

N215422 001 Nov 22, 2021

+

10MG/PACKET

N215422 002 Nov 22, 2021

+!

20MG/PACKET

N215422 003 Nov 22, 2021

PRESCRIPTION DRUG PRODUCT LIST

BACLOFEN

INJECTABLE; INTRATHECAL

BACLOFEN

AP	AMNEAL	<u>0.05MG/ML</u>	<u>A091193</u>	<u>001</u>	May 03, 2016
AP		<u>0.5MG/ML</u>	<u>A091193</u>	<u>002</u>	May 03, 2016
AP		<u>2MG/ML</u>	<u>A091193</u>	<u>003</u>	May 03, 2016
AP	MAIA PHARMS INC	<u>0.05MG/ML</u>	<u>A210777</u>	<u>001</u>	Jan 15, 2021
AP		<u>0.5MG/ML</u>	<u>A210048</u>	<u>001</u>	Sep 11, 2019
AP		<u>1MG/ML</u>	<u>A210315</u>	<u>001</u>	Jul 30, 2019
AP		<u>2MG/ML</u>	<u>A210048</u>	<u>002</u>	Sep 11, 2019
AP	MYLAN LABS LTD	<u>0.5MG/ML</u>	<u>A209592</u>	<u>001</u>	Mar 21, 2018
AP		<u>1MG/ML</u>	<u>A209594</u>	<u>001</u>	Mar 06, 2018
AP		<u>2MG/ML</u>	<u>A209592</u>	<u>002</u>	Mar 21, 2018
AP	RUBICON	<u>0.5MG/ML</u>	<u>A217324</u>	<u>001</u>	Feb 22, 2023
AP		<u>1MG/ML</u>	<u>A217324</u>	<u>002</u>	Feb 22, 2023
AP		<u>2MG/ML</u>	<u>A217324</u>	<u>003</u>	Feb 22, 2023

GABLOFEN

AP	+! PIRAMAL CRITICAL	<u>0.05MG/ML</u>	<u>N022462</u>	<u>001</u>	Nov 19, 2010
AP	+!	<u>0.5MG/ML</u>	<u>N022462</u>	<u>002</u>	Nov 19, 2010
AP	+!	<u>1MG/ML</u>	<u>N022462</u>	<u>004</u>	Jun 22, 2012
AP	+!	<u>2MG/ML</u>	<u>N022462</u>	<u>003</u>	Nov 19, 2010

LIORESAL

AP	+! AMNEAL	<u>0.05MG/ML</u>	<u>N020075</u>	<u>003</u>	Nov 07, 1996
AP	+!	<u>0.5MG/ML</u>	<u>N020075</u>	<u>001</u>	Jun 17, 1992
AP	+!	<u>2MG/ML</u>	<u>N020075</u>	<u>002</u>	Jun 17, 1992

SOLUTION; ORAL

BACLOFEN

	RUBICON	5MG/5ML	A214445	001	Dec 06, 2024
	OZOBAX DS				
	+! METACEL PHARMS LLC	10MG/5ML	N208193	002	Oct 12, 2023

SUSPENSION; ORAL

BACLOFEN

AB	ANI PHARMS	<u>25MG/5ML</u>	<u>A217252</u>	<u>001</u>	Jun 08, 2023
AB	+! AZURITY	<u>25MG/5ML</u>	<u>N215602</u>	<u>001</u>	Feb 04, 2022

TABLET; ORAL

BACLOFEN

AB	ANDAS 5 HOLDING	<u>5MG</u>	<u>A212378</u>	<u>003</u>	Apr 30, 2021
AB		<u>10MG</u>	<u>A212378</u>	<u>001</u>	Oct 09, 2020
AB		<u>20MG</u>	<u>A212378</u>	<u>002</u>	Oct 09, 2020
AB	AUROBINDO PHARMA LTD	<u>10MG</u>	<u>A214099</u>	<u>001</u>	Jul 13, 2021
AB		<u>20MG</u>	<u>A214099</u>	<u>002</u>	Jul 13, 2021
AB	BEXIMCO PHARMS USA	<u>5MG</u>	<u>A214114</u>	<u>003</u>	Sep 19, 2023
AB		<u>10MG</u>	<u>A214114</u>	<u>001</u>	Jul 16, 2021
AB		<u>20MG</u>	<u>A214114</u>	<u>002</u>	Jul 16, 2021
AB	ENDO OPERATIONS	<u>10MG</u>	<u>A077068</u>	<u>002</u>	Aug 30, 2005
AB		<u>20MG</u>	<u>A077068</u>	<u>001</u>	Aug 30, 2005
AB	GRAVITI PHARMS	<u>5MG</u>	<u>A217788</u>	<u>003</u>	Oct 24, 2024
AB		<u>10MG</u>	<u>A217788</u>	<u>001</u>	Jan 10, 2024
AB		<u>20MG</u>	<u>A217788</u>	<u>002</u>	Jan 10, 2024
AB	IVAX SUB TEVA PHARMS	<u>10MG</u>	<u>A072234</u>	<u>001</u>	Jul 21, 1988
AB	!	<u>20MG</u>	<u>A072235</u>	<u>001</u>	Jul 21, 1988
AB	LANNETT CO INC	<u>5MG</u>	<u>A077241</u>	<u>003</u>	Sep 22, 2021
AB		<u>10MG</u>	<u>A077241</u>	<u>002</u>	Jul 06, 2007
AB		<u>20MG</u>	<u>A077241</u>	<u>001</u>	Dec 20, 2005
AB	MANKIND PHARMA	<u>5MG</u>	<u>A215885</u>	<u>001</u>	Jan 25, 2022
AB		<u>10MG</u>	<u>A215885</u>	<u>002</u>	Jan 25, 2022
AB		<u>20MG</u>	<u>A215885</u>	<u>003</u>	Jan 25, 2022
AB	MICRO LABS	<u>5MG</u>	<u>A217687</u>	<u>001</u>	Nov 08, 2023
AB		<u>10MG</u>	<u>A217687</u>	<u>002</u>	Nov 08, 2023
AB		<u>20MG</u>	<u>A217687</u>	<u>003</u>	Nov 08, 2023
AB	NORTHSTAR HLTHCARE	<u>10MG</u>	<u>A078401</u>	<u>002</u>	Sep 18, 2009
AB		<u>20MG</u>	<u>A078401</u>	<u>001</u>	Sep 18, 2009
AB	OXFORD PHARMS	<u>10MG</u>	<u>A077088</u>	<u>002</u>	Oct 31, 2007
AB		<u>20MG</u>	<u>A077088</u>	<u>001</u>	Oct 31, 2007
AB	RISING	<u>5MG</u>	<u>A214374</u>	<u>001</u>	Mar 05, 2021
AB		<u>10MG</u>	<u>A214374</u>	<u>002</u>	Mar 05, 2021
AB		<u>20MG</u>	<u>A214374</u>	<u>003</u>	Mar 05, 2021
AB	!	<u>5MG</u>	<u>A209102</u>	<u>001</u>	Nov 28, 2017
AB		<u>10MG</u>	<u>A209102</u>	<u>002</u>	Nov 28, 2017
AB		<u>15MG</u>	<u>A209102</u>	<u>004</u>	Feb 05, 2024

PRESCRIPTION DRUG PRODUCT LISTBACLOFEN

TABLET; ORAL

BACLOFEN

AB		20MG	A209102 003	Nov 28, 2017
AB	UNICHEM	5MG	A212067 003	Mar 09, 2023
AB		10MG	A212067 001	Jul 09, 2020
AB		20MG	A212067 002	Jul 09, 2020
AB	UPSHER SMITH LABS	10MG	A074584 001	Aug 19, 1996
AB		20MG	A074584 002	Aug 19, 1996
AB	ZYDUS LIFESCIENCES	5MG	A211659 003	Apr 17, 2020
AB		10MG	A211659 001	Nov 23, 2018
AB		15MG	A211659 004	Feb 13, 2024
AB		20MG	A211659 002	Nov 23, 2018
BX	HIBROW HLTHCARE	5MG	A211555 003	Nov 30, 2021
BX		10MG	A211555 001	Feb 01, 2019
BX		20MG	A211555 002	Feb 01, 2019

BALOXAVIR MARBOXIL

FOR SUSPENSION; ORAL

XOFLUZA

+! GENENTECH INC 2MG/ML N214410 001 Nov 23, 2020

TABLET; ORAL

XOFLUZA

+ GENENTECH INC 40MG N210854 002 Oct 24, 2018

+! 80MG N210854 003 Mar 18, 2021

BALSALAZIDE DISODIUM

CAPSULE; ORAL

BALSALAZIDE DISODIUM

AB	APOTEX INC	750MG	A077883 001	Dec 28, 2007
AB	HIKMA	750MG	A077806 001	Dec 28, 2007
AB	ZYDUS LIFESCIENCES	750MG	A217592 001	Jun 08, 2023

COLAZAL

AB	+! VALEANT PHARMS INTL	750MG	N020610 001	Jul 18, 2000
-----------	------------------------	--------------	--------------------	--------------

BARICITINIB

TABLET; ORAL

OLUMIANT

+ ELI LILLY AND CO 1MG N207924 002 Oct 08, 2019

+ 2MG N207924 001 May 31, 2018

+! 4MG N207924 003 May 10, 2022

BARIUM SULFATE

FOR SUSPENSION; ORAL

E-Z-HD

+! BRACCO 98% (334GM/BOT) N208036 001 Jan 11, 2016

E-Z-PAQUE

+! BRACCO 96% (169GM/BOT) N208036 002 Apr 07, 2017

VARIBAR THIN LIQUID

+! BRACCO 81% (120GM/BOT) N208036 004 Apr 30, 2019

PASTE; ORAL

VARIBAR PUDDING

BRACCO 40% N208844 001 Oct 14, 2016

SUSPENSION; ORAL

ENTERO VU 24%

+! BRACCO 24% (144GM/600ML) N208143 008 May 29, 2020

LIQUID E-Z-PAQUE

+! BRACCO 60% (213GM/BOT) N208143 003 Mar 01, 2017

READI-CAT 2

+! BRACCO 2% (9GM/BOT) N208143 001 Jan 15, 2016

READI-CAT 2 SMOOTHIE

+! BRACCO 2% (9GM/BOT) N208143 002 Jan 15, 2016

TAGITOL V

+! BRACCO 40% (8GM/BOT) N208143 005 Aug 04, 2017

VARIBAR HONEY

+! BRACCO 40% (100GM/250ML) N208143 007 Mar 26, 2018

VARIBAR NECTAR

+! BRACCO 40% (96GM/240ML) N208143 004 Jul 07, 2017

VARIBAR THIN HONEY

+! BRACCO 40% (100GM/250ML) N208143 006 Jan 23, 2018

PRESCRIPTION DRUG PRODUCT LIST

BAZEDOXIFENE ACETATE; ESTROGENS, CONJUGATED

TABLET; ORAL

DUAVEE

+! WYETH PHARMS EQ 20MG BASE;0.45MG N022247 001 Oct 03, 2013

BECLOMETHASONE DIPROPIONATE

AEROSOL, METERED; INHALATION

QVAR REDIHALER

+ NORTON WATERFORD 0.04MG/INH N207921 001 Aug 03, 2017

+ 0.08MG/INH N207921 002 Aug 03, 2017

AEROSOL, METERED; NASAL

QNASL

+ TEVA BRANDED PHARM 0.04MG/ACTUATION N202813 002 Dec 17, 2014

+! 0.08MG/ACTUATION N202813 001 Mar 23, 2012

BEDAQUILINE FUMARATE

TABLET; ORAL

SIRTURO

+ JANSSEN THERAP EQ 20MG BASE N204384 002 May 27, 2020

+! EQ 100MG BASE N204384 001 Dec 28, 2012

BELINOSTAT

POWDER; INTRAVENOUS

BELEODAQ

+! ACROTECH BIOPHARMA 500MG/VIAL N206256 001 Jul 03, 2014

BELUMOSUDIL MESYLATE

TABLET; ORAL

REZUROCK

+! KADMON PHARMS LLC EQ 200MG BASE N214783 001 Jul 16, 2021

BELZUTIFAN

TABLET; ORAL

WELIREG

+! MERCK SHARP DOHME 40MG N215383 001 Aug 13, 2021

BEMPEDOIC ACID

TABLET; ORAL

NEXLETOL

+! ESPERION THERAPS 180MG N211616 001 Feb 21, 2020

INC

BEMPEDOIC ACID; EZETIMIBE

TABLET; ORAL

NEXLIZET

+! ESPERION THERAPS 180MG;10MG N211617 001 Feb 26, 2020

INC

BENAZEPRIL HYDROCHLORIDE

TABLET; ORAL

BENAZEPRIL HYDROCHLORIDE

AB	AMNEAL PHARMS	5MG	A076820 001	Feb 03, 2006
AB		10MG	A076820 002	Feb 03, 2006
AB		20MG	A076820 003	Feb 03, 2006
AB		40MG	A076820 004	Feb 03, 2006
AB	ANI PHARMS	5MG	A076333 001	Feb 11, 2004
AB		10MG	A076333 002	Feb 11, 2004
AB		20MG	A076333 003	Feb 11, 2004
AB		40MG	A076333 004	Feb 11, 2004
AB	AUROBINDO PHARMA	10MG	A078212 001	May 22, 2008
AB		20MG	A078212 002	May 22, 2008
AB	!	40MG	A078212 003	May 22, 2008
AB	CHARTWELL RX	5MG	A076402 001	Feb 11, 2004
AB		10MG	A076402 002	Feb 11, 2004
AB		20MG	A076402 003	Feb 11, 2004
AB		40MG	A076402 004	Feb 11, 2004
AB	COREPHARMA	5MG	A077128 001	Mar 08, 2006
AB		10MG	A077128 002	Mar 08, 2006
AB		20MG	A077128 003	Mar 08, 2006
AB		40MG	A077128 004	Mar 08, 2006
AB	HERITAGE PHARMA	5MG	A076267 001	Feb 11, 2004
AB		10MG	A076267 002	Feb 11, 2004
AB		20MG	A076267 003	Feb 11, 2004
AB		40MG	A076267 004	Feb 11, 2004
AB	PRINSTON INC	5MG	A076118 001	Feb 11, 2004
AB		10MG	A076118 002	Feb 11, 2004
AB		20MG	A076118 003	Feb 11, 2004

PRESCRIPTION DRUG PRODUCT LIST

BENAZEPRIL HYDROCHLORIDE

TABLET; ORAL

BENAZEPRIL HYDROCHLORIDE

<u>AB</u>		<u>40MG</u>	<u>A076118 004</u>	Feb 11, 2004
<u>AB</u>	SUN PHARM INDS LTD	<u>5MG</u>	<u>A076344 001</u>	Feb 11, 2004
<u>AB</u>		<u>10MG</u>	<u>A076344 002</u>	Feb 11, 2004
<u>AB</u>		<u>20MG</u>	<u>A076344 003</u>	Feb 11, 2004
<u>AB</u>		<u>40MG</u>	<u>A076344 004</u>	Feb 11, 2004
<u>AB</u>	TEVA	<u>5MG</u>	<u>A076211 001</u>	Feb 11, 2004
<u>AB</u>		<u>10MG</u>	<u>A076211 002</u>	Feb 11, 2004
<u>AB</u>		<u>20MG</u>	<u>A076211 003</u>	Feb 11, 2004
<u>AB</u>		<u>40MG</u>	<u>A076211 004</u>	Feb 11, 2004
<u>AB</u>	ZYDUS LIFESCIENCES	<u>5MG</u>	<u>A078848 001</u>	May 23, 2008
<u>AB</u>		<u>10MG</u>	<u>A078848 002</u>	May 23, 2008
<u>AB</u>		<u>20MG</u>	<u>A078848 003</u>	May 23, 2008
<u>AB</u>		<u>40MG</u>	<u>A078848 004</u>	May 23, 2008

LOTENSIN

<u>AB</u>	+	VALIDUS PHARMS	<u>10MG</u>	<u>N019851 002</u>	Jun 25, 1991
<u>AB</u>	+		<u>20MG</u>	<u>N019851 003</u>	Jun 25, 1991
<u>AB</u>	+		<u>40MG</u>	<u>N019851 004</u>	Jun 25, 1991

BENAZEPRIL HYDROCHLORIDE; HYDROCHLOROTHIAZIDE

TABLET; ORAL

BENAZEPRIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE

<u>AB</u>	ANI PHARMS	<u>5MG; 6.25MG</u>	<u>A076342 001</u>	Feb 11, 2004
<u>AB</u>		<u>10MG; 12.5MG</u>	<u>A076342 002</u>	Feb 11, 2004
<u>AB</u>		<u>20MG; 12.5MG</u>	<u>A076342 003</u>	Feb 11, 2004
<u>AB</u>		<u>20MG; 25MG</u>	<u>A076342 004</u>	Feb 11, 2004
<u>AB</u>	APOTEX	<u>5MG; 6.25MG</u>	<u>A078794 001</u>	Aug 21, 2014
<u>AB</u>		<u>10MG; 12.5MG</u>	<u>A078794 002</u>	Aug 21, 2014
<u>AB</u>		<u>20MG; 12.5MG</u>	<u>A078794 003</u>	Aug 21, 2014
<u>AB</u>		<u>20MG; 25MG</u>	<u>A078794 004</u>	Aug 21, 2014
<u>AB</u>	SANDOZ	<u>5MG; 6.25MG</u>	<u>A076631 001</u>	Feb 11, 2004
<u>AB</u>		<u>10MG; 12.5MG</u>	<u>A076631 002</u>	Feb 11, 2004
<u>AB</u>		<u>20MG; 12.5MG</u>	<u>A076631 003</u>	Feb 11, 2004
<u>AB</u>		<u>20MG; 25MG</u>	<u>A076631 004</u>	Feb 11, 2004

LOTENSIN HCT

<u>AB</u>	+	VALIDUS PHARMS	<u>10MG; 12.5MG</u>	<u>N020033 002</u>	May 19, 1992
<u>AB</u>	+		<u>20MG; 12.5MG</u>	<u>N020033 004</u>	May 19, 1992
<u>AB</u>	+		<u>20MG; 25MG</u>	<u>N020033 003</u>	May 19, 1992

BENDAMUSTINE HYDROCHLORIDE

POWDER; INTRAVENOUS

BENDAMUSTINE HYDROCHLORIDE

<u>AP</u>	ACCORD HLTHCARE	<u>25MG/VIAL</u>	<u>A205574 001</u>	Dec 07, 2022
<u>AP</u>		<u>100MG/VIAL</u>	<u>A205574 002</u>	Dec 07, 2022
<u>AP</u>	APOTEX	<u>25MG/VIAL</u>	<u>A204230 001</u>	Jun 05, 2023
<u>AP</u>		<u>100MG/VIAL</u>	<u>A204230 002</u>	Jun 05, 2023
<u>AP</u>	DR REDDYS	<u>25MG/VIAL</u>	<u>A205376 001</u>	Dec 07, 2022
<u>AP</u>		<u>100MG/VIAL</u>	<u>A205376 002</u>	Dec 07, 2022
<u>AP</u>	EUGIA PHARMA	<u>25MG/VIAL</u>	<u>A214739 001</u>	Jun 05, 2023
<u>AP</u>		<u>100MG/VIAL</u>	<u>A214739 002</u>	Jun 05, 2023
<u>AP</u>	MEITHEAL	<u>25MG/VIAL</u>	<u>A211001 001</u>	Jun 05, 2023
<u>AP</u>		<u>100MG/VIAL</u>	<u>A211001 002</u>	Jun 05, 2023

TREANDA

<u>AP</u>	+	CEPHALON	<u>25MG/VIAL</u>	<u>N022249 002</u>	May 01, 2009
<u>AP</u>	+		<u>100MG/VIAL</u>	<u>N022249 001</u>	Mar 20, 2008

SOLUTION; INTRAVENOUS

BELRAPZO

<u>AP</u>	+	EAGLE PHARMS	<u>100MG/4ML (25MG/ML)</u>	<u>N205580 001</u>	May 15, 2018
-----------	---	--------------	----------------------------	--------------------	--------------

BENDAMUSTINE HYDROCHLORIDE

<u>AP</u>	+	APOTEX	<u>100MG/4ML (25MG/ML)</u>	<u>N215033 001</u>	Dec 07, 2022
<u>AP</u>	+	BAXTER HLTHCARE CORP	<u>100MG/4ML (25MG/ML)</u>	<u>N216078 001</u>	Dec 15, 2022

BENDEKA

	+	EAGLE PHARMS	100MG/4ML (25MG/ML)	N208194 001	Dec 07, 2015
--	---	--------------	---------------------	-------------	--------------

VIVIMUSTA

	+	AZURITY	100MG/4ML (25MG/ML)	N212209 001	Dec 07, 2022
--	---	---------	---------------------	-------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

BENOXINATE HYDROCHLORIDE; FLUORESCEIN SODIUM

SOLUTION/DROPS;OPHTHALMIC

ALTAFLUOR BENOX

+! ALTAIRE PHARMS INC 0.4%;0.25%

N208582 001 Dec 14, 2017

FLUORESCEIN SODIUM AND BENOXINATE HYDROCHLORIDE

+! BAUSCH LOMB IRELAND 0.4%;0.3%

N211039 001 Mar 09, 2020

BENZGALANTAMINE GLUCONATE

TABLET, DELAYED RELEASE;ORAL

ZUNVEYL

+! ALPHA COGNITION EQ 5MG BASE

N218549 001 Jul 26, 2024

+ EQ 10MG BASE

N218549 002 Jul 26, 2024

+ EQ 15MG BASE

N218549 003 Jul 26, 2024

BENZNIDAZOLE

TABLET;ORAL

BENZNIDAZOLE

+ CHEMO RESEARCH SL 12.5MG

N209570 001 Aug 29, 2017

+! 100MG

N209570 002 Aug 29, 2017

BENZONATATE

CAPSULE;ORAL

BENZONATATEAA ASCENT PHARMS INC100MGA211518 001 Feb 22, 2019AA 200MGA211518 003 Feb 22, 2019AA BIONPHARMA 100MGA081297 001 Jan 29, 1993AA 200MGA081297 002 Oct 30, 2007AA CSPEC-NBP PHARM 100MGA202765 002 Aug 25, 2017AA 200MGA202765 001 Jul 31, 2015AA HERITAGE PHARMS 100MGA040682 001 Jul 30, 2007

LABS

AA 200MGA040682 002 Jul 30, 2007AA ! PURACAP PHARM LLC 100MGA206948 001 Dec 19, 2018AA ! 200MGA206948 002 Dec 19, 2018AA STRIDES SOFTGELS 100MGA091133 001 Jul 30, 2015AA 200MGA091133 002 Jul 30, 2015AA THEPHARMANETWORK 100MGA040627 001 Mar 30, 2007

LLC

AA 200MGA040627 002 Jul 25, 2007AA ZYDUS PHARMS USA 100MGA040597 001 Jun 08, 2007AA 200MGA040597 002 Jun 08, 2007TESSALONAA + PFIZER 100MGN011210 001

BENZONATATE

! THEPHARMANETWORK 150MG

A040627 003 Sep 24, 2014

LLC

BENZOYL PEROXIDE

CREAM;TOPICAL

EPSOLAY

+! GALDERMA LABS LP 5%

N214510 001 Apr 22, 2022

BENZOYL PEROXIDE; CLINDAMYCIN PHOSPHATE

GEL;TOPICAL

ACANYAAB +! BAUSCH 2.5%;EQ 1.2% BASEN050819 001 Oct 23, 2008CLINDAMYCIN PHOSPHATE AND BENZOYL PEROXIDEAB ACTAVIS LABS UT INC 2.5%;EQ 1.2% BASEA205128 001 Jun 19, 2015AB ENCUBE 2.5%;EQ 1.2% BASEA207194 001 Aug 19, 2019AB 5%;1.2%A212433 001 Apr 28, 2021AB ! GLENMARK SPECLT 5%;EQ 1% BASEA209252 001 Mar 14, 2019AB MYLAN PHARMS INC 5%;EQ 1% BASEA065443 001 Aug 11, 2009AB PADAGIS ISRAEL 2.5%;EQ 1.2% BASEA205397 001 Sep 09, 2019AB 3.75%;EQ 1.2% BASEA209610 001 Aug 20, 2024AB 5%;EQ 1% BASEA202440 001 Sep 21, 2015AB ! 5%;1.2%A090979 001 Jun 26, 2012AB TARO 2.5%;EQ 1.2% BASEA206575 001 Aug 19, 2019AB 3.75%;EQ 1.2% BASEA208683 001 Jun 05, 2018AB 5%;EQ 1% BASEA208776 001 May 25, 2018AB 5%;1.2%A206218 001 Dec 15, 2017AB ZYDUS PHARMS 5%;1.2%A210794 001 Dec 28, 2018DUACAB + STIEFEL 5%;1.2%N050741 001 Aug 26, 2002ONEXTONAB +! BAUSCH 3.75%;EQ 1.2% BASEN050819 002 Nov 24, 2014

PRESCRIPTION DRUG PRODUCT LIST

BENZOYL PEROXIDE; ERYTHROMYCIN

GEL; TOPICAL

BENZAMYCIN

AB	+ !	VALEANT INTL	5%;3%	N050557 001	Oct 26, 1984
-----------	------------	--------------	--------------	--------------------	--------------

ERYTHROMYCIN AND BENZOYL PEROXIDE

AB		RISING	5%;3%	A065385 001	Sep 18, 2015
-----------	--	--------	--------------	--------------------	--------------

BENZOYL PEROXIDE; TRETINOIN

CREAM; TOPICAL

TWYNEO

+ !	GALDERMA LABS LP	3%;0.1%	N214902 001	Jul 26, 2021
------------	------------------	----------------	--------------------	--------------

BENZPHETAMINE HYDROCHLORIDE

TABLET; ORAL

BENZPHETAMINE HYDROCHLORIDE

AA		CHARTWELL	50MG	A090473 002	Sep 15, 2010
-----------	--	-----------	-------------	--------------------	--------------

AA		EPIC PHARMA LLC	50MG	A090346 001	Dec 15, 2015
-----------	--	-----------------	-------------	--------------------	--------------

AA	!	KVK TECH	50MG	A090968 001	Jul 20, 2010
-----------	----------	----------	-------------	--------------------	--------------

		CHARTWELL	25MG	A090473 001	Sep 15, 2010
--	--	-----------	-------------	--------------------	--------------

BENZTROPINE MESYLATE

INJECTABLE; INJECTION

BENZTROPINE MESYLATE

AP		FRESENIUS KABI USA	1MG/ML	A090233 001	Jul 28, 2009
-----------	--	--------------------	---------------	--------------------	--------------

AP		HIKMA	1MG/ML	A209442 001	Oct 14, 2021
-----------	--	-------	---------------	--------------------	--------------

AP	!	HIKMA FARMACEUTICA	1MG/ML	A090287 001	Aug 31, 2009
-----------	----------	--------------------	---------------	--------------------	--------------

AP		NAVINTA LLC	1MG/ML	A091525 001	Feb 05, 2013
-----------	--	-------------	---------------	--------------------	--------------

TABLET; ORAL

BENZTROPINE MESYLATE

AA		AIPING PHARM INC	0.5MG	A040103 001	Dec 12, 1996
-----------	--	------------------	--------------	--------------------	--------------

AA			1MG	A040103 002	Dec 12, 1996
-----------	--	--	------------	--------------------	--------------

AA			2MG	A040103 003	Dec 12, 1996
-----------	--	--	------------	--------------------	--------------

AA		CHARTWELL RX	0.5MG	A081265 003	Nov 29, 2023
-----------	--	--------------	--------------	--------------------	--------------

AA			1MG	A081265 002	Jan 23, 1992
-----------	--	--	------------	--------------------	--------------

AA			2MG	A081265 001	Jan 23, 1992
-----------	--	--	------------	--------------------	--------------

AA		ENDO OPERATIONS	0.5MG	A040715 001	Aug 27, 2007
-----------	--	-----------------	--------------	--------------------	--------------

AA			1MG	A040715 002	Aug 27, 2007
-----------	--	--	------------	--------------------	--------------

AA			2MG	A040715 003	Aug 27, 2007
-----------	--	--	------------	--------------------	--------------

AA		EPIC PHARMA LLC	0.5MG	A072264 001	Feb 27, 1989
-----------	--	-----------------	--------------	--------------------	--------------

AA			1MG	A072265 001	Feb 27, 1989
-----------	--	--	------------	--------------------	--------------

AA			2MG	A072266 001	Feb 27, 1989
-----------	--	--	------------	--------------------	--------------

AA		LEADING	0.5MG	A090168 001	Nov 28, 2012
-----------	--	---------	--------------	--------------------	--------------

AA			1MG	A090168 002	Nov 28, 2012
-----------	--	--	------------	--------------------	--------------

AA			2MG	A090168 003	Nov 28, 2012
-----------	--	--	------------	--------------------	--------------

AA	!	NUVO PHARM	0.5MG	A204713 001	Apr 14, 2015
-----------	----------	------------	--------------	--------------------	--------------

AA	!		1MG	A204713 002	Apr 14, 2015
-----------	----------	--	------------	--------------------	--------------

AA	!		2MG	A204713 003	Apr 14, 2015
-----------	----------	--	------------	--------------------	--------------

AA		PLIVA	0.5MG	A089058 001	Aug 10, 1988
-----------	--	-------	--------------	--------------------	--------------

AA			1MG	A089059 001	Aug 10, 1988
-----------	--	--	------------	--------------------	--------------

AA			2MG	A089060 001	Aug 10, 1988
-----------	--	--	------------	--------------------	--------------

BEPOTASTINE BESILATE

SOLUTION/DROPS; OPHTHALMIC

BEPOTASTINE BESILATE

AT		ALEMBIC	1.5%	A214588 001	Apr 05, 2023
-----------	--	---------	-------------	--------------------	--------------

AT		APOTEX	1.5%	A206066 001	Mar 05, 2019
-----------	--	--------	-------------	--------------------	--------------

AT		MYLAN	1.5%	A206220 001	Mar 18, 2019
-----------	--	-------	-------------	--------------------	--------------

AT		SOMERSET THERAPS LLC	1.5%	A217770 001	Aug 14, 2024
-----------	--	-------------------------	-------------	--------------------	--------------

BEPREVE

AT	+ !	BAUSCH AND LOMB INC	1.5%	N022288 001	Sep 08, 2009
-----------	------------	---------------------	-------------	--------------------	--------------

BERDAZIMER SODIUM

GEL; TOPICAL

ZELSUVMI

+ !	LNHC	EQ 10.3% BASE	N217424 001	Jan 05, 2024
------------	------	----------------------	--------------------	--------------

BEROTRALSTAT HYDROCHLORIDE

CAPSULE; ORAL

ORLADEYO

+	BIOCRIST	EQ 110MG BASE	N214094 001	Dec 03, 2020
----------	----------	----------------------	--------------------	--------------

+ !		EQ 150MG BASE	N214094 002	Dec 03, 2020
------------	--	----------------------	--------------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

BESIFLOXACIN HYDROCHLORIDE

SUSPENSION/DROPS;OPHTHALMIC

BESIVANCE

+! BAUSCH AND LOMB EQ 0.6% BASE N022308 001 May 28, 2009

BETAINE

FOR SOLUTION;ORAL

BETAINE

AB ETON 1GM/SCOOPFUL A210508 001 Jan 28, 2022

AB NOVITIUM PHARMA 1GM/SCOOPFUL A214864 001 Nov 23, 2021

CYSTADANE

AB +! RECORDATI RARE 1GM/SCOOPFUL N020576 001 Oct 25, 1996

BETAMETHASONE ACETATE; BETAMETHASONE SODIUM PHOSPHATE

INJECTABLE; INJECTION

BETAMETHASONE ACETATE AND BETAMETHASONE SODIUM PHOSPHATE

AB AM REGENT 3MG/ML;EQ 3MG BASE/ML A090747 001 Jul 31, 2009

AB HIKMA 3MG/ML;EQ 3MG BASE/ML A077838 001 Jan 17, 2023

CELESTONE SOLUSPAN

AB +! ORGANON 3MG/ML;EQ 3MG BASE/ML N014602 001

BETAMETHASONE DIPROPIONATE

CREAM; TOPICAL

BETAMETHASONE DIPROPIONATE

AB ACTAVIS MID EQ 0.05% BASE A070885 001 Feb 03, 1987

ATLANTIC

AB COSETTE EQ 0.05% BASE A210217 001 Oct 12, 2018

AB +! FOUGERA PHARMS EQ 0.05% BASE N019137 001 Jun 26, 1984

AB TARO EQ 0.05% BASE A073552 001 Apr 30, 1992

AB ZYDUS PHARMS EQ 0.05% BASE A208885 001 Jan 11, 2019

CREAM, AUGMENTED; TOPICAL

BETAMETHASONE DIPROPIONATE

AB FOUGERA PHARMS EQ 0.05% BASE A076215 001 Dec 09, 2003

AB ! GLENMARK PHARMS INC EQ 0.05% BASE A078930 001 Sep 23, 2008

AB PADAGIS ISRAEL EQ 0.05% BASE A076592 001 Dec 09, 2003

AB TARO EQ 0.05% BASE A076543 001 Dec 09, 2003

GEL, AUGMENTED; TOPICAL

BETAMETHASONE DIPROPIONATE

AB ! FOUGERA PHARMS EQ 0.05% BASE A075276 001 May 13, 2003

AB TARO EQ 0.05% BASE A076508 001 Dec 02, 2003

LOTION; TOPICAL

BETAMETHASONE DIPROPIONATE

AB COSETTE EQ 0.05% BASE A071467 001 Aug 10, 1987

AB ! FOUGERA PHARMS INC EQ 0.05% BASE A070275 001 Aug 12, 1985

AB PADAGIS US EQ 0.05% BASE A072538 001 Jan 31, 1990

LOTION, AUGMENTED; TOPICAL

BETAMETHASONE DIPROPIONATE

AB FOUGERA PHARMS EQ 0.05% BASE A077111 001 May 21, 2007

AB ! TARO EQ 0.05% BASE A077477 001 May 21, 2007

OINTMENT; TOPICAL

BETAMETHASONE DIPROPIONATE

AB ACTAVIS MID EQ 0.05% BASE A071012 001 Feb 03, 1987

ATLANTIC

AB +! FOUGERA PHARMS INC EQ 0.05% BASE N019141 001 Sep 04, 1984

AB PADAGIS ISRAEL EQ 0.05% BASE A215847 001 Apr 12, 2022

AB TARO EQ 0.05% BASE A074271 001 Sep 15, 1994

AB TRUPHARMA EQ 0.05% BASE A215186 001 Feb 18, 2022

AB ZYDUS LIFESCIENCES EQ 0.05% BASE A214048 001 Jul 14, 2020

OINTMENT, AUGMENTED; TOPICAL

BETAMETHASONE DIPROPIONATE

AB ACTAVIS MID EQ 0.05% BASE A074304 001 Aug 31, 1995

ATLANTIC

AB AUROBINDO PHARMA EQ 0.05% BASE A218289 001 Apr 10, 2024

LTD

AB FOUGERA PHARMS EQ 0.05% BASE A075373 001 Jun 22, 1999

AB LUPIN LTD EQ 0.05% BASE A209106 001 Dec 18, 2019

AB TARO EQ 0.05% BASE A076753 001 Oct 12, 2004

DIPROLENE

AB +! ORGANON EQ 0.05% BASE N018741 001 Jul 27, 1983

SPRAY; TOPICAL

SERNIVO

+! PRIMUS PHARMS EQ 0.05% BASE/SPRAY N208079 001 Feb 05, 2016

PRESCRIPTION DRUG PRODUCT LIST

BETAMETHASONE DIPROPIONATE; CALCIPOTRIENE

AEROSOL, FOAM;TOPICAL

ENSTILAR

+! LEO PHARMA AS 0.064%;0.005% N207589 001 Oct 16, 2015

CREAM;TOPICAL

WYNZORA

+! MC2 0.064%;0.005% N213422 001 Jul 20, 2020

OINTMENT;TOPICAL

CALCIPOTRIENE AND BETAMETHASONE DIPROPIONATE**AB** PADAGIS ISRAEL 0.064%;0.005% **A200174 001** Dec 12, 2014TACLONEX**AB** +! LEO PHARMA AS 0.064%;0.005% **N021852 001** Jan 09, 2006

SUSPENSION;TOPICAL

CALCIPOTRIENE AND BETAMETHASONE DIPROPIONATE**AB** COSETTE 0.064%;0.005% **A210765 001** May 11, 2020**AB** TARO 0.064%;0.005% **A213269 001** Sep 02, 2020CALCIPOTRIENE AND BETHAMETHASONE DIPROPIONATE**AB** PADAGIS ISRAEL 0.064%;0.005% **A212367 001** Sep 11, 2020TACLONEX**AB** +! LEO PHARMA AS 0.064%;0.005% **N022185 001** May 09, 2008BETAMETHASONE DIPROPIONATE; CLOTRIMAZOLE

CREAM;TOPICAL

CLOTRIMAZOLE AND BETAMETHASONE DIPROPIONATE**AB** ACTAVIS MID EQ 0.05% BASE;1% **A076002 001** Aug 02, 2002

ATLANTIC

AB FOUGERA PHARMS EQ 0.05% BASE;1% **A075502 001** Jun 05, 2001**AB** GLENMARK SPECLT EQ 0.05% BASE;1% **A202894 001** Oct 30, 2015**AB** ! TARO EQ 0.05% BASE;1% **A075673 001** May 29, 2001

LOTION;TOPICAL

CLOTRIMAZOLE AND BETAMETHASONE DIPROPIONATE**AB** FOUGERA PHARMS EQ 0.05% BASE;1% **A076516 001** Jun 16, 2005**AB** ! TARO EQ 0.05% BASE;1% **A076493 001** Jul 28, 2004BETAMETHASONE VALERATE

AEROSOL, FOAM;TOPICAL

BETAMETHASONE VALERATE**AB** ALEMBIC 0.12% **A215832 001** Aug 22, 2024**AB** PADAGIS ISRAEL 0.12% **A078337 001** Nov 26, 2012**AB** ! TARO 0.12% **A208204 001** May 24, 2017**AB** XIROMED 0.12% **A210639 001** Apr 18, 2023

CREAM;TOPICAL

BETA-VAL**AB** COSETTE EQ 0.1% BASE **N018642 001** Mar 24, 1983BETAMETHASONE VALERATE**AB** +! FOUGERA PHARMS INC EQ 0.1% BASE **N018861 001** Aug 31, 1983DERMABET**AB** TARO EQ 0.1% BASE **A072041 001** Jan 06, 1988VALNAC**AB** ACTAVIS MID EQ 0.1% BASE **A070050 001** Oct 10, 1984

ATLANTIC

LOTION;TOPICAL

BETAMETHASONE VALERATE**AB** +! FOUGERA PHARMS INC EQ 0.1% BASE **N018866 001** Aug 31, 1983**AB** SCIEGEN PHARMS INC EQ 0.1% BASE **A070052 001** Jul 31, 1985

OINTMENT;TOPICAL

BETA-VAL**AB** COSETTE EQ 0.1% BASE **A070069 001** Dec 19, 1985BETAMETHASONE VALERATE**AB** ACTAVIS MID EQ 0.1% BASE **A070051 001** Oct 10, 1984

ATLANTIC

AB +! FOUGERA PHARMS INC EQ 0.1% BASE **N018865 001** Aug 31, 1983BETAXOLOL HYDROCHLORIDE

SOLUTION/DROPS;OPHTHALMIC

BETAXOLOL HYDROCHLORIDE**AT** ACELLA EQ 0.5% BASE **A078694 001** Nov 16, 2009**AT** MEDIMETRIKS PHARMS EQ 0.5% BASE **A075630 001** Apr 12, 2001BETOPTIC**AT** +! SANDOZ EQ 0.5% BASE **N019270 001** Aug 30, 1985

SUSPENSION/DROPS;OPHTHALMIC

BETOPTIC S

+! NOVARTIS EQ 0.25% BASE N019845 001 Dec 29, 1989

PRESCRIPTION DRUG PRODUCT LIST

BETAXOLOL HYDROCHLORIDE

TABLET; ORAL

BETAXOLOL HYDROCHLORIDE

<u>AB</u>		EPIC PHARMA	<u>10MG</u>	<u>A075541</u>	<u>001</u>	Oct 22, 1999
<u>AB</u>	!		<u>20MG</u>	<u>A075541</u>	<u>002</u>	Oct 22, 1999
<u>AB</u>		KVK TECH	<u>10MG</u>	<u>A078962</u>	<u>001</u>	Jun 27, 2008
<u>AB</u>			<u>20MG</u>	<u>A078962</u>	<u>002</u>	Jun 27, 2008

BETHANECHOL CHLORIDE

TABLET; ORAL

BETHANECHOL CHLORIDE

<u>AA</u>	!	AMNEAL PHARM	<u>5MG</u>	<u>A040855</u>	<u>001</u>	Nov 21, 2007
<u>AA</u>	!		<u>10MG</u>	<u>A040855</u>	<u>002</u>	Nov 21, 2007
<u>AA</u>	!		<u>25MG</u>	<u>A040855</u>	<u>003</u>	Nov 21, 2007
<u>AA</u>	!		<u>50MG</u>	<u>A040855</u>	<u>004</u>	Nov 21, 2007
<u>AA</u>		CHARTWELL RX	<u>5MG</u>	<u>A040728</u>	<u>002</u>	Oct 26, 2007
<u>AA</u>			<u>10MG</u>	<u>A040728</u>	<u>003</u>	Oct 26, 2007
<u>AA</u>			<u>25MG</u>	<u>A040728</u>	<u>004</u>	Oct 26, 2007
<u>AA</u>			<u>50MG</u>	<u>A040728</u>	<u>001</u>	Oct 26, 2007
<u>AA</u>		UPSHER SMITH LABS	<u>5MG</u>	<u>A040633</u>	<u>001</u>	Jun 01, 2005
<u>AA</u>			<u>10MG</u>	<u>A040634</u>	<u>001</u>	Jun 01, 2005
<u>AA</u>			<u>25MG</u>	<u>A040635</u>	<u>001</u>	Jun 01, 2005
<u>AA</u>			<u>50MG</u>	<u>A040636</u>	<u>001</u>	Jun 01, 2005

DUVOID

<u>AA</u>		CHARTWELL RX	<u>10MG</u>	<u>A086262</u>	<u>001</u>	
<u>AA</u>			<u>25MG</u>	<u>A086263</u>	<u>001</u>	
<u>AA</u>			<u>50MG</u>	<u>A085882</u>	<u>003</u>	

BEXAGLIFLOZIN

TABLET; ORAL

BRENZAVVY

	+	!	THERACOSBIO	20MG	N214373	001	Jan 20, 2023
--	---	---	-------------	------	---------	-----	--------------

BEXAROTENE

CAPSULE; ORAL

BEXAROTENE

<u>AB</u>		AMNEAL PHARMS NY	<u>75MG</u>	<u>A210105</u>	<u>001</u>	Sep 04, 2018
<u>AB</u>		ANI PHARMS	<u>75MG</u>	<u>A209861</u>	<u>001</u>	May 08, 2018
<u>AB</u>		ASCENT PHARMS INC	<u>75MG</u>	<u>A208628</u>	<u>001</u>	Sep 06, 2024
<u>AB</u>		BIONPHARMA	<u>75MG</u>	<u>A203174</u>	<u>001</u>	Aug 12, 2014
<u>AB</u>		CIPLA	<u>75MG</u>	<u>A210352</u>	<u>001</u>	Dec 10, 2024
<u>AB</u>		HIKMA	<u>75MG</u>	<u>A203663</u>	<u>001</u>	Jun 16, 2020
<u>AB</u>		TEVA PHARMS USA	<u>75MG</u>	<u>A209931</u>	<u>001</u>	Jan 14, 2021
<u>AB</u>		UPSHER SMITH LABS	<u>75MG</u>	<u>A209886</u>	<u>001</u>	Jul 25, 2018

TARGRETIN

<u>AB</u>	+	!	VALEANT LUXEMBOURG	<u>75MG</u>	<u>N021055</u>	<u>001</u>	Dec 29, 1999
-----------	---	---	--------------------	-------------	----------------	------------	--------------

GEL; TOPICAL

BEXAROTENE

<u>AB</u>		AMNEAL	<u>1%</u>	<u>A215398</u>	<u>001</u>	Apr 27, 2022
-----------	--	--------	-----------	----------------	------------	--------------

TARGRETIN

<u>AB</u>	+	!	BAUSCH	<u>1%</u>	<u>N021056</u>	<u>001</u>	Jun 28, 2000
-----------	---	---	--------	-----------	----------------	------------	--------------

BICALUTAMIDE

TABLET; ORAL

BICALUTAMIDE

<u>AB</u>		ACCORD HLTHCARE	<u>50MG</u>	<u>A078917</u>	<u>001</u>	Jul 06, 2009
<u>AB</u>		ADAPTIS	<u>50MG</u>	<u>A079089</u>	<u>001</u>	Jul 06, 2009
<u>AB</u>		APOTEX	<u>50MG</u>	<u>A200274</u>	<u>001</u>	May 21, 2015
<u>AB</u>		CHARTWELL RX	<u>50MG</u>	<u>A091011</u>	<u>001</u>	Jun 10, 2015
<u>AB</u>		SANDOZ	<u>50MG</u>	<u>A078575</u>	<u>001</u>	Jul 06, 2009
<u>AB</u>		SUN PHARM	<u>50MG</u>	<u>A079110</u>	<u>001</u>	Jul 06, 2009
<u>AB</u>		WATSON LABS TEVA	<u>50MG</u>	<u>A078634</u>	<u>001</u>	Aug 28, 2009

CASODEX

<u>AB</u>	+	!	ANI PHARMS	<u>50MG</u>	<u>N020498</u>	<u>001</u>	Oct 04, 1995
-----------	---	---	------------	-------------	----------------	------------	--------------

BICTEGRAVIR SODIUM; EMTRICITABINE; TENOFOVIR ALAFENAMIDE FUMARATE

TABLET; ORAL

BIKTARVY

	+		GILEAD SCIENCES INC	EQ 30MG BASE;120MG;EQ 15MG BASE	N210251	002	Oct 07, 2021
	+	!		EQ 50MG BASE;200MG;EQ 25MG BASE	N210251	001	Feb 07, 2018

PRESCRIPTION DRUG PRODUCT LIST

BIMATOPROST

IMPLANT;OPHTHALMIC

DURYSTA

+! ABBVIE

10MCG

N211911 001 Mar 04, 2020

SOLUTION/DROPS;OPHTHALMIC

BIMATOPROST

AT	!	ALEMBIC	0.03%	A210263	001	Apr 12, 2019
AT		APOTEX	0.03%	A090449	001	Jul 20, 2015
AT		EUGIA PHARMA	0.03%	A205537	001	Oct 06, 2022
AT		GLAND PHARMA LTD	0.03%	A210126	001	Mar 22, 2019
AT		LUPIN LTD	0.03%	A203991	001	Feb 20, 2015
AT		MICRO LABS	0.03%	A202505	001	Sep 08, 2020
AT		SANDOZ	0.03%	A202565	001	May 05, 2015
AT		SOMERSET THERAPS LLC	0.03%	A207601	001	Jun 19, 2019

LUMIGAN

+! ABBVIE

0.01%

N022184 001 Aug 31, 2010

SOLUTION/DROPS;TOPICAL

BIMATOPROST

AT		ALEMBIC	0.03%	A210515	001	Jan 21, 2020
AT		APOTEX	0.03%	A201894	001	Dec 01, 2014
AT		SANDOZ	0.03%	A202719	001	Apr 19, 2016

LATISSE

AT	+	!	ABBVIE	0.03%	N022369	001	Dec 24, 2008
-----------	---	---	--------	--------------	----------------	------------	--------------

BINIMETINIB

TABLET;ORAL

MEKTOVI

+! ARRAY BIOPHARMA INC

15MG

N210498 001 Jun 27, 2018

BIRCH TRITERPENES

GEL;TOPICAL

FILSUVEZ

+! CHIESI

10%

N215064 001 Dec 18, 2023

BISMUTH SUBCITRATE POTASSIUM; METRONIDAZOLE; TETRACYCLINE HYDROCHLORIDE

CAPSULE;ORAL

BISMUTH SUBCITRATE POTASSIUM, METRONIDAZOLE AND TETRACYCLINE HYDROCHLORIDE

AB		ENDO OPERATIONS	140MG;125MG;125MG	A205770	001	Mar 06, 2023	
AB		NOVAST LABS	140MG;125MG;125MG	A217511	001	Jul 03, 2023	
AB	+	!	LABS JUVISE	140MG;125MG;125MG	N050786	001	Sep 28, 2006

BISMUTH SUBSALICYLATE; METRONIDAZOLE; TETRACYCLINE HYDROCHLORIDE

TABLET, CHEWABLE, TABLET, CAPSULE;ORAL

BISMUTH SUBSALICYLATE, METRONIDAZOLE AND TETRACYCLINE HYDROCHLORIDE

! AILEX PHARMS LLC 262.4MG, N/A, N/A;N/A, 250MG, N/A;N/A,
N/A, 500MG

A202584 001 Nov 30, 2018

BISOPROLOL FUMARATE

TABLET;ORAL

BISOPROLOL FUMARATE

AB		ALEMBIC	5MG	A204891	001	Jan 11, 2017
AB			10MG	A204891	002	Jan 11, 2017
AB		AUROBINDO PHARMA	5MG	A077910	001	Dec 27, 2006
AB			10MG	A077910	002	Dec 27, 2006
AB		HARMAN FINOCHEM	5MG	A217617	001	Jan 18, 2024
AB			10MG	A217617	002	Jan 18, 2024
AB		NOVITIUM PHARMA	5MG	A215563	001	Oct 29, 2021
AB			10MG	A215563	002	Oct 29, 2021
AB		PRINSTON INC	5MG	A217368	001	Jul 14, 2023
AB			10MG	A217368	002	Jul 14, 2023
AB		RUBICON	5MG	A075643	001	Nov 16, 2000
AB			10MG	A075643	002	Nov 16, 2000
AB		TRUPHARMA	5MG	A075474	001	Oct 25, 2002
AB			10MG	A075474	002	Oct 25, 2002
AB		UNICHEM	5MG	A078635	001	Aug 18, 2009
AB	!		10MG	A078635	002	Aug 18, 2009
AB		ZYDUS LIFESCIENCES	5MG	A215680	001	Jul 25, 2022
AB			10MG	A215680	002	Jul 25, 2022

PRESCRIPTION DRUG PRODUCT LIST

BISOPROLOL FUMARATE; HYDROCHLOROTHIAZIDE

TABLET; ORAL

BISOPROLOL FUMARATE AND HYDROCHLOROTHIAZIDE

<u>AB</u>	CADILA	<u>2.5MG; 6.25MG</u>	<u>A215666 001</u>	Nov 04, 2022
<u>AB</u>		<u>5MG; 6.25MG</u>	<u>A215666 002</u>	Nov 04, 2022
<u>AB</u>		<u>10MG; 6.25MG</u>	<u>A215666 003</u>	Nov 04, 2022
<u>AB</u>	EDENBRIDGE PHARMS	<u>2.5MG; 6.25MG</u>	<u>A212678 001</u>	Jul 09, 2020
<u>AB</u>		<u>5MG; 6.25MG</u>	<u>A212678 002</u>	Jul 09, 2020
<u>AB</u>		<u>10MG; 6.25MG</u>	<u>A212678 003</u>	Jul 09, 2020
<u>AB</u>	EPIC PHARMA LLC	<u>2.5MG; 6.25MG</u>	<u>A075579 001</u>	Sep 25, 2000
<u>AB</u>		<u>5MG; 6.25MG</u>	<u>A075579 002</u>	Sep 25, 2000
<u>AB</u>		<u>10MG; 6.25MG</u>	<u>A075579 003</u>	Sep 25, 2000
<u>AB</u>	GLENMARK PHARMS LTD	<u>2.5MG; 6.25MG</u>	<u>A215995 001</u>	Jan 26, 2022
<u>AB</u>		<u>5MG; 6.25MG</u>	<u>A215995 002</u>	Jan 26, 2022
<u>AB</u>		<u>10MG; 6.25MG</u>	<u>A215995 003</u>	Jan 26, 2022
<u>AB</u>	MYLAN	<u>2.5MG; 6.25MG</u>	<u>A075768 001</u>	Sep 25, 2000
<u>AB</u>		<u>5MG; 6.25MG</u>	<u>A075768 002</u>	Sep 25, 2000
<u>AB</u>		<u>10MG; 6.25MG</u>	<u>A075768 003</u>	Sep 25, 2000
<u>AB</u>	NOVITIUM PHARMA	<u>2.5MG; 6.25MG</u>	<u>A215562 001</u>	Nov 04, 2021
<u>AB</u>		<u>5MG; 6.25MG</u>	<u>A215562 002</u>	Nov 04, 2021
<u>AB</u>		<u>10MG; 6.25MG</u>	<u>A215562 003</u>	Nov 04, 2021
<u>AB</u>	UNICHEM	<u>2.5MG; 6.25MG</u>	<u>A079106 001</u>	Jul 28, 2010
<u>AB</u>		<u>5MG; 6.25MG</u>	<u>A079106 002</u>	Jul 28, 2010
<u>AB</u>		<u>10MG; 6.25MG</u>	<u>A079106 003</u>	Jul 28, 2010
<u>ZIAC</u>				
<u>AB</u>	+ TEVA BRANDED PHARM	<u>2.5MG; 6.25MG</u>	<u>N020186 003</u>	Mar 26, 1993
<u>AB</u>	+	<u>5MG; 6.25MG</u>	<u>N020186 001</u>	Mar 26, 1993
<u>AB</u>	+!	<u>10MG; 6.25MG</u>	<u>N020186 002</u>	Mar 26, 1993

BIVALIRUDIN

INJECTABLE; INTRAVENOUS

ANGIOMAX

<u>AP</u>	+! SANDOZ	<u>250MG/VIAL</u>	<u>N020873 001</u>	Dec 15, 2000
-----------	-----------	-------------------	--------------------	--------------

BIVALIRUDIN

<u>AP</u>	ACCORD HLTHCARE	<u>250MG/VIAL</u>	<u>A206551 001</u>	Nov 22, 2017
<u>AP</u>	DR REDDYS	<u>250MG/VIAL</u>	<u>A201577 001</u>	May 26, 2017
<u>AP</u>	EUGIA PHARMA	<u>250MG/VIAL</u>	<u>A205962 001</u>	Jul 27, 2018
<u>AP</u>	FRESENIUS KABI USA	<u>250MG/VIAL</u>	<u>A090189 001</u>	Oct 28, 2016
<u>AP</u>	MEITHEAL	<u>250MG/VIAL</u>	<u>A091602 001</u>	Jul 16, 2018
<u>AP</u>	MYLAN INSTITUTIONAL	<u>250MG/VIAL</u>	<u>A202471 001</u>	Jun 01, 2018
<u>AP</u>	SHUANGCHENG	<u>250MG/VIAL</u>	<u>A210031 001</u>	Oct 23, 2019
<u>AP</u>	SLATE RUN PHARMA	<u>250MG/VIAL</u>	<u>A213078 001</u>	May 28, 2021

SOLUTION; INTRAVENOUS

ANGIOMAX RTU

	+! MAIA PHARMS INC	250MG/50ML (5MG/ML)	N211215 001	Jul 25, 2019
--	--------------------	---------------------	-------------	--------------

BLEOMYCIN SULFATE

INJECTABLE; INJECTION

BLEOMYCIN SULFATE

<u>AP</u>	! FRESENIUS KABI USA	<u>EQ 15 UNITS BASE/VIAL</u>	<u>A065185 001</u>	Jan 28, 2008
<u>AP</u>	!	<u>EQ 30 UNITS BASE/VIAL</u>	<u>A065185 002</u>	Jan 28, 2008
<u>AP</u>	HIKMA	<u>EQ 15 UNITS BASE/VIAL</u>	<u>A065042 002</u>	Oct 17, 2001
<u>AP</u>		<u>EQ 30 UNITS BASE/VIAL</u>	<u>A065042 001</u>	Oct 17, 2001
<u>AP</u>	HOSPIRA	<u>EQ 15 UNITS BASE/VIAL</u>	<u>A065031 001</u>	Mar 10, 2000
<u>AP</u>		<u>EQ 30 UNITS BASE/VIAL</u>	<u>A065031 002</u>	Mar 10, 2000
<u>AP</u>	MEITHEAL	<u>EQ 15 UNITS BASE/VIAL</u>	<u>A205030 001</u>	Apr 20, 2018
<u>AP</u>		<u>EQ 30 UNITS BASE/VIAL</u>	<u>A205030 002</u>	Apr 20, 2018

BORTEZOMIB

INJECTABLE; INTRAVENOUS, SUBCUTANEOUS

BORTEZOMIB

<u>AP</u>	APOTEX	<u>3.5MG/VIAL</u>	<u>A205533 001</u>	May 02, 2022
<u>AP</u>	BAXTER HLTHCARE CORP	<u>3.5MG/VIAL</u>	<u>A213823 001</u>	May 02, 2022
<u>AP</u>	DR REDDYS	<u>3.5MG/VIAL</u>	<u>A202963 001</u>	Jul 26, 2022
<u>AP</u>	EUGIA PHARMA	<u>3.5MG/VIAL</u>	<u>A212825 001</u>	May 02, 2022
<u>AP</u>	FRESENIUS KABI USA	<u>3.5MG/VIAL</u>	<u>A209659 001</u>	May 02, 2022
<u>AP</u>	HETERO LABS LTD VI	<u>3.5MG/VIAL</u>	<u>A212204 001</u>	May 03, 2024
<u>AP</u>	HOSPIRA	<u>3.5MG/VIAL</u>	<u>A208460 001</u>	Jul 26, 2022
<u>AP</u>	JIANGSU HANSOH PHARM	<u>3.5MG/VIAL</u>	<u>A215011 001</u>	Jul 26, 2022
<u>AP</u>	MEITHEAL	<u>3.5MG/VIAL</u>	<u>A212958 001</u>	Jul 26, 2022
<u>AP</u>	MSN	<u>3.5MG/VIAL</u>	<u>A209622 001</u>	Jul 26, 2022
<u>AP</u>	PHARMSCIENCE INC	<u>3.5MG/VIAL</u>	<u>A208392 001</u>	May 02, 2022

PRESCRIPTION DRUG PRODUCT LIST

BORTEZOMIB

INJECTABLE; INTRAVENOUS, SUBCUTANEOUS

BORTEZOMIB

AP	QILU PHARM HAINAN	3.5MG/VIAL	A210824 001	May 02, 2022
AP	RELIANCE LIFE SCI	3.5MG/VIAL	A211898 001	Oct 11, 2022
AP	SANDOZ	3.5MG/VIAL	A203654 001	Jul 26, 2022
AP	ZYDUS PHARMS	3.5MG/VIAL	A210204 001	May 02, 2022

VELCADE

AP	+ TAKEDA PHARMS USA	3.5MG/VIAL	N021602 001	May 13, 2003
POWDER; INTRAVENOUS, SUBCUTANEOUS				
BORTEZOMIB				
	+ HOSPIRA	1MG/VIAL	N209191 001	May 02, 2022
	+	2.5MG/VIAL	N209191 002	May 02, 2022
SOLUTION; INTRAVENOUS, SUBCUTANEOUS				
BORTEZOMIB				
	+ SHILPA	3.5MG/1.4ML (2.5MG/ML)	N212782 001	Aug 26, 2024

BOSENTAN

TABLET; ORAL

BOSENTAN

AB	SUN PHARM	62.5MG	A209324 001	Apr 26, 2019
AB		125MG	A209324 002	Apr 26, 2019
AB	WATSON LABS INC	62.5MG	A207110 001	Apr 26, 2019
AB		125MG	A207110 002	Apr 26, 2019
AB	ZYDUS PHARMS	62.5MG	A207760 001	Apr 26, 2019
AB		125MG	A207760 002	Apr 26, 2019

TRACLEER

AB	+ ACTELION	62.5MG	N021290 001	Nov 20, 2001
AB	+	125MG	N021290 002	Nov 20, 2001
TABLET, FOR SUSPENSION; ORAL				
TRACLEER				
	+ ACTELION	32MG	N209279 001	Sep 05, 2017

BOSUTINIB MONOHYDRATE

CAPSULE; ORAL

BOSULIF

	+ PF PRISM CV	EQ 50MG BASE	N217729 001	Sep 26, 2023
	+	EQ 100MG BASE	N217729 002	Sep 26, 2023
TABLET; ORAL				
BOSULIF				
	+ PF PRISM CV	EQ 100MG BASE	N203341 001	Sep 04, 2012
	+	EQ 400MG BASE	N203341 003	Oct 27, 2017
	+	EQ 500MG BASE	N203341 002	Sep 04, 2012

BREMELANOTIDE ACETATE

SOLUTION; SUBCUTANEOUS

VYLEESI (AUTOINJECTOR)

	+ COSETTE	EQ 1.75MG BASE/0.3ML (EQ 1.75MG BASE/0.3 ML)	N210557 001	Jun 21, 2019
--	------------------	--	-------------	--------------

BREXANOLONE

SOLUTION; INTRAVENOUS

ZULRESSO

	+ SAGE THERAP	100MG/20ML (5MG/ML)	N211371 001	Jun 17, 2019
--	----------------------	---------------------	-------------	--------------

BREXPIRAZOLE

TABLET; ORAL

REXULTI

	+ OTSUKA	0.25MG	N205422 001	Jul 10, 2015
	+	0.5MG	N205422 002	Jul 10, 2015
	+	1MG	N205422 003	Jul 10, 2015
	+	2MG	N205422 004	Jul 10, 2015
	+	3MG	N205422 005	Jul 10, 2015
	+	4MG	N205422 006	Jul 10, 2015

BRIGATINIB

TABLET; ORAL

ALUNBRIG

	+ TAKEDA PHARMS USA	30MG	N208772 001	Apr 28, 2017
	+	90MG	N208772 002	Apr 28, 2017
	+	180MG	N208772 003	Oct 02, 2017

PRESCRIPTION DRUG PRODUCT LIST

BRILLIANT BLUE GSOLUTION;OPHTHALMIC
TISSUEBLUE

+! DUTCH OPTHALMIC 0.025% N209569 001 Dec 20, 2019

BRIMONIDINE TARTRATE

GEL;TOPICAL

BRIMONIDINE TARTRATEAB PADAGIS ISRAEL EQ 0.33% BASE A209158 001 Sep 23, 2021MIRVASOAB +! GALDERMA LABS LP EQ 0.33% BASE N204708 001 Aug 23, 2013

SOLUTION/DROPS;OPHTHALMIC

ALPHAGAN PAB +! ABBVIE 0.1% N021770 001 Aug 19, 2005BRIMONIDINE TARTRATEAB ALEMBIC 0.1% A216909 001 Aug 01, 2023AB APOTEX 0.1% A078480 001 Dec 21, 2022AB LUPIN LTD 0.1% A213215 001 Aug 26, 2024AB SANDOZ 0.1% A203172 001 Mar 04, 2024AB SOMERSET THERAPS 0.1% A216906 001 Aug 09, 2024

LLC

ALPHAGAN PAT +! ABBVIE 0.15% N021262 001 Mar 16, 2001BRIMONIDINE TARTRATEAT ALEMBIC 0.15% A215225 001 Mar 29, 2023AT APOTEX 0.15% A078479 001 Jan 31, 2022AT ! BAUSCH AND LOMB 0.2% A076260 001 May 28, 2003AT INDOCO 0.2% A091691 001 Nov 18, 2014AT MICRO LABS 0.2% A217846 001 Nov 22, 2023AT RISING 0.2% A076439 001 Mar 14, 2006AT SANDOZ 0.2% A076254 001 Sep 16, 2003AT 0.2% A078075 001 Jan 30, 2008AT SOMERSET THERAPS 0.2% A208992 001 Mar 11, 2019

LLC

OOLIANAAT +! SANDOZ 0.15% N021764 001 May 22, 2006BRIMONIDINE TARTRATE; BRINZOLAMIDESUSPENSION/DROPS;OPHTHALMIC
SIMBRINZA

+! ALCON LABS INC 0.2%;1% N204251 001 Apr 19, 2013

BRIMONIDINE TARTRATE; TIMOLOL MALEATE

SOLUTION/DROPS;OPHTHALMIC

BRIMONIDINE TARTRATE AND TIMOLOL MALEATEAB ALEMBIC 0.2%;EQ 0.5% BASE A215230 001 Aug 25, 2023AB AMNEAL 0.2%;EQ 0.5% BASE A217288 001 Dec 30, 2024AB APOTEX 0.2%;EQ 0.5% BASE A091442 001 Apr 20, 2022AB FLORIDA 0.2%;EQ 0.5% BASE A201949 001 Oct 04, 2022AB GLENMARK PHARMS LTD 0.2%;EQ 0.5% BASE A214987 001 May 16, 2024AB SANDOZ 0.2%;EQ 0.5% BASE A091087 001 Apr 04, 2022AB SENTISS 0.2%;EQ 0.5% BASE A091086 001 Oct 31, 2022AB SOMERSET 0.2%;EQ 0.5% BASE A216114 001 Sep 10, 2024AB UPSHER SMITH LABS 0.2%;EQ 0.5% BASE A215598 001 Dec 12, 2022COMBIGANAB +! ABBVIE 0.2%;EQ 0.5% BASE N021398 001 Oct 30, 2007BRINCIDOFIVIR

SUSPENSION;ORAL

TEMBEXA

+! EMERGENT BIODEFENSE 10MG/ML N214460 001 Jun 04, 2021

TABLET;ORAL

TEMBEXA

+! EMERGENT BIODEFENSE 100MG N214461 001 Jun 04, 2021

BRINZOLAMIDE

SUSPENSION/DROPS;OPHTHALMIC

AZOPTAB +! SANDOZ 1% N020816 001 Apr 01, 1998BRINZOLAMIDEAB BAUSCH AND LOMB 1% A204884 001 Aug 18, 2021AB PADAGIS US 1% A211914 001 Jul 28, 2023AB WATSON LABS INC 1% A209406 001 Nov 27, 2020

PRESCRIPTION DRUG PRODUCT LIST

BRIVARACETAM

SOLUTION; INTRAVENOUS

BRIVIACT

+! UCB INC 50MG/5ML (10MG/ML) N205837 001 May 12, 2016

SOLUTION; ORAL

BRIVIACT

+! UCB INC 10MG/ML N205838 001 May 12, 2016

TABLET; ORAL

BRIVIACT

+ UCB INC 10MG N205836 001 May 12, 2016

+ 25MG N205836 002 May 12, 2016

+ 50MG N205836 003 May 12, 2016

+ 75MG N205836 004 May 12, 2016

+! 100MG N205836 005 May 12, 2016

BROMFENAC SODIUM

SOLUTION/DROPS; OPHTHALMIC

BROMFENAC SODIUMAB ALEMBIC EQ 0.07% ACID A214340 001 Jul 08, 2024AB AMNEAL EQ 0.07% ACID A210962 001 Jul 29, 2024AB APOTEX EQ 0.07% ACID A207334 001 Jul 08, 2024AB LUPIN LTD EQ 0.07% ACID A206027 001 Nov 22, 2023AB EQ 0.075% ACID A211239 001 Feb 02, 2024BROMSITEAB +! SUN PHARM EQ 0.075% ACID N206911 001 Apr 08, 2016PROLENSAAB +! BAUSCH AND LOMB EQ 0.07% ACID N203168 001 Apr 05, 2013BROMFENAC SODIUMAT2 ! ALEMBIC EQ 0.09% ACID A210560 001 Jun 21, 2019AT2 EUGIA PHARMA EQ 0.09% ACID A204813 001 Mar 18, 2022AT2 GLAND PHARMA LTD EQ 0.09% ACID A211029 001 Mar 17, 2020AT2 LUPIN LTD EQ 0.09% ACID A202903 001 Aug 15, 2023AT2 SENTISS EQ 0.09% ACID A203395 001 Jan 22, 2014BROMOCRIPTINE MESYLATE

CAPSULE; ORAL

BROMOCRIPTINE MESYLATEAB ! MYLAN EQ 5MG BASE A077226 001 Apr 04, 2005AB ZYDUS PHARMS USA EQ 5MG BASE A078899 001 Jul 30, 2008PARLODELAB + ESJAY PHARMA EQ 5MG BASE N017962 002 Mar 01, 1982

TABLET; ORAL

BROMOCRIPTINE MESYLATEAB ! PADAGIS US EQ 2.5MG BASE A077646 001 Oct 01, 2008AB SANDOZ EQ 2.5MG BASE A074631 001 Jan 13, 1998PARLODELAB + ESJAY PHARMA EQ 2.5MG BASE N017962 001

CYCLOSET

+! VEROSCIENCE EQ 0.8MG BASE N020866 001 May 05, 2009

BROMPHENIRAMINE MALEATE; DEXTROMETHORPHAN HYDROBROMIDE; PSEUDOEPHEDRINE HYDROCHLORIDE

SYRUP; ORAL

BROMFED-DMAA + WOCKHARDT BIO AG 2MG/5ML; 10MG/5ML; 30MG/5ML A088811 001 Jun 07, 1985BROMPHENIRAMINE MALEATE, PSEUDOEPHEDRINE HYDROCHLORIDE AND DEXTROMETHORPHAN HYDROBROMIDEAA ACELLA 2MG/5ML; 10MG/5ML; 30MG/5ML A203375 001 Sep 20, 2016AA ALKEM LABS LTD 2MG/5ML; 10MG/5ML; 30MG/5ML A210647 001 Jul 14, 2020AA CHARTWELL MOLECULAR 2MG/5ML; 10MG/5ML; 30MG/5ML A213125 001 Apr 17, 2020AA DR REDDYS LABS SA 2MG/5ML; 10MG/5ML; 30MG/5ML A207676 001 Dec 04, 2018AA ! PADAGIS US 2MG/5ML; 10MG/5ML; 30MG/5ML A205292 001 Jul 15, 2014AA PHARM ASSOC 2MG/5ML; 10MG/5ML; 30MG/5ML A202940 001 Jul 21, 2014AA TARO 2MG/5ML; 10MG/5ML; 30MG/5ML A205112 001 Feb 27, 2017AA WES PHARMA INC 2MG/5ML; 10MG/5ML; 30MG/5ML A211170 001 Jun 16, 2020BUDESONIDE

AEROSOL, FOAM; RECTAL

BUDESONIDEAB ! PADAGIS ISRAEL 2MG/ACTUATION A215328 001 Apr 12, 2023UCERISAB + SALIX 2MG/ACTUATION N205613 001 Oct 07, 2014

CAPSULE, DELAYED RELEASE; ORAL

BUDESONIDEAB AMNEAL PHARMS 3MG A206200 001 Jul 31, 2017AB AUROBINDO PHARMA 3MG A090410 001 May 16, 2011

PRESCRIPTION DRUG PRODUCT LIST

BUDESONIDE

CAPSULE, DELAYED RELEASE;ORAL

BUDESONIDE

USA

<u>AB</u>	DR REDDYS LABS SA	<u>3MG</u>	<u>A206623</u>	<u>001</u>	Apr 08, 2016
<u>AB</u>	RISING	<u>3MG</u>	<u>A207367</u>	<u>001</u>	Apr 07, 2017
<u>AB</u>	ZYDUS PHARMS	<u>3MG</u>	<u>A206134</u>	<u>001</u>	May 04, 2017

ENTOCORT EC

<u>AB</u>	+!	PADAGIS US	<u>3MG</u>	<u>N021324</u>	<u>001</u>	Oct 02, 2001
		TARPEYO				
		+!	CALLIDITAS	4MG	N215935	001 Dec 15, 2021

POWDER, METERED;INHALATION

PULMICORT FLEXHALER

		+!	CHEPLAPHARM	0.08MG/INH	N021949	001 Jul 12, 2006
			+!		N021949	002 Jul 12, 2006

SUSPENSION; INHALATION

BUDESONIDE

<u>AN</u>		CIPLA	<u>0.25MG/2ML</u>	<u>A205710</u>	<u>001</u>	Nov 16, 2017
<u>AN</u>			<u>0.5MG/2ML</u>	<u>A205710</u>	<u>002</u>	Nov 16, 2017
<u>AN</u>			<u>1MG/2ML</u>	<u>A205710</u>	<u>003</u>	Nov 16, 2017
<u>AN</u>		IMPAX LABS INC	<u>0.25MG/2ML</u>	<u>A078404</u>	<u>001</u>	Jul 31, 2012
<u>AN</u>			<u>0.5MG/2ML</u>	<u>A078404</u>	<u>002</u>	Jul 31, 2012
<u>AN</u>		LUPIN	<u>0.5MG/2ML</u>	<u>A210897</u>	<u>001</u>	Nov 09, 2018
<u>AN</u>		NEPHRON	<u>0.25MG/2ML</u>	<u>A078202</u>	<u>001</u>	Mar 30, 2009
<u>AN</u>			<u>0.5MG/2ML</u>	<u>A078202</u>	<u>002</u>	Mar 30, 2009
<u>AN</u>		SANDOZ	<u>0.25MG/2ML</u>	<u>A201966</u>	<u>003</u>	Sep 27, 2013
<u>AN</u>			<u>0.5MG/2ML</u>	<u>A201966</u>	<u>002</u>	Sep 27, 2013
<u>AN</u>			<u>1MG/2ML</u>	<u>A201966</u>	<u>001</u>	Sep 27, 2013
<u>AN</u>		SUN PHARM	<u>0.25MG/2ML</u>	<u>A211922</u>	<u>001</u>	Apr 14, 2021
<u>AN</u>			<u>0.5MG/2ML</u>	<u>A211922</u>	<u>002</u>	Apr 14, 2021
<u>AN</u>			<u>1MG/2ML</u>	<u>A211922</u>	<u>003</u>	Apr 14, 2021
<u>AN</u>		TEVA PHARMS	<u>0.25MG/2ML</u>	<u>A077519</u>	<u>001</u>	Nov 18, 2008
<u>AN</u>			<u>0.5MG/2ML</u>	<u>A077519</u>	<u>002</u>	Nov 18, 2008
<u>AN</u>		TEVA PHARMS USA	<u>1MG/2ML</u>	<u>A204548</u>	<u>001</u>	Mar 08, 2016

PULMICORT RESPULES

<u>AN</u>		+!	ASTRAZENECA	<u>0.25MG/2ML</u>	<u>N020929</u>	<u>001</u>	Aug 08, 2000
<u>AN</u>			+!		<u>N020929</u>	<u>002</u>	Aug 08, 2000
<u>AN</u>			+!		<u>N020929</u>	<u>003</u>	Aug 08, 2000

SUSPENSION;ORAL

EOHILIA

		+!	TAKEDA PHARMS USA	2MG/10ML	N213976	001	Feb 09, 2024
--	--	----	-------------------	----------	---------	-----	--------------

TABLET, EXTENDED RELEASE;ORAL

BUDESONIDE

<u>AB</u>		ACTAVIS LABS FL INC	<u>9MG</u>	<u>A205457</u>	<u>001</u>	Jul 03, 2018
<u>AB</u>		MYLAN	<u>9MG</u>	<u>A208851</u>	<u>001</u>	Sep 17, 2020

UCERIS

<u>AB</u>		+!	SALIX	<u>9MG</u>	<u>N203634</u>	<u>001</u>	Jan 14, 2013
-----------	--	----	-------	------------	----------------	------------	--------------

BUDESONIDE; FORMOTEROL FUMARATE

AEROSOL, METERED;INHALATION

SYMBICORT AEROSPHERE

		+!	ASTRAZENECA	0.16MG/INH;0.0048MG/INH	N216579	001	Apr 28, 2023
--	--	----	-------------	-------------------------	---------	-----	--------------

BUDESONIDE; FORMOTEROL FUMARATE DIHYDRATE

AEROSOL, METERED;INHALATION

BREYNA

<u>AB</u>		MYLAN	<u>0.08MG/INH;0.0045MG/INH</u>	<u>A211699</u>	<u>001</u>	Mar 15, 2022
<u>AB</u>			<u>0.16MG/INH;0.0045MG/INH</u>	<u>A211699</u>	<u>002</u>	Mar 15, 2022

SYMBICORT

<u>AB</u>		+!	ASTRAZENECA	<u>0.08MG/INH;0.0045MG/INH</u>	<u>N021929</u>	<u>001</u>	Jul 21, 2006
<u>AB</u>			+!		<u>N021929</u>	<u>002</u>	Jul 21, 2006

BUDESONIDE; FORMOTEROL FUMARATE; GLYCOPYRROLATE

AEROSOL, METERED;INHALATION

BREZTRI AEROSPHERE

		+!	ASTRAZENECA AB	0.16MG/INH;0.0048MG/INH;0.009MG/INH	N212122	001	Jul 23, 2020
--	--	----	----------------	-------------------------------------	---------	-----	--------------

BUMETANIDE

INJECTABLE; INJECTION

BUMETANIDE

<u>AP</u>		GLAND PHARMA LTD	<u>0.25MG/ML</u>	<u>A216434</u>	<u>001</u>	May 26, 2022
<u>AP</u>		LUPIN LTD	<u>0.25MG/ML</u>	<u>A217153</u>	<u>001</u>	Sep 17, 2024
<u>AP</u>		MSN	<u>0.25MG/ML</u>	<u>A215364</u>	<u>001</u>	Aug 04, 2022
<u>AP</u>		QILU PHARM HAINAN	<u>0.25MG/ML</u>	<u>A219116</u>	<u>001</u>	Sep 19, 2024

PRESCRIPTION DRUG PRODUCT LIST

BUMETANIDE

INJECTABLE; INJECTION

BUMETANIDE

<u>AP</u>	SAGENT	<u>0.25MG/ML</u>	<u>A074441</u>	<u>001</u>	Jan 27, 1995
<u>AP</u>	! WEST-WARD PHARMS INT	<u>0.25MG/ML</u>	<u>A079196</u>	<u>001</u>	Apr 30, 2008

TABLET; ORAL

BUMETANIDE

<u>AB</u>	AMNEAL PHARMS CO	<u>0.5MG</u>	<u>A209724</u>	<u>001</u>	Oct 18, 2017
<u>AB</u>		<u>1MG</u>	<u>A209724</u>	<u>002</u>	Oct 18, 2017
<u>AB</u>		<u>2MG</u>	<u>A209724</u>	<u>003</u>	Oct 18, 2017
<u>AB</u>	APPCO	<u>0.5MG</u>	<u>A212931</u>	<u>001</u>	Sep 18, 2024
<u>AB</u>		<u>1MG</u>	<u>A212931</u>	<u>002</u>	Sep 18, 2024
<u>AB</u>		<u>2MG</u>	<u>A212931</u>	<u>003</u>	Sep 18, 2024
<u>AB</u>	HERITAGE PHARMA	<u>0.5MG</u>	<u>A074225</u>	<u>001</u>	Apr 24, 1995
<u>AB</u>		<u>1MG</u>	<u>A074225</u>	<u>002</u>	Apr 24, 1995
<u>AB</u>		<u>2MG</u>	<u>A074225</u>	<u>003</u>	Apr 24, 1995
<u>AB</u>	RISING	<u>0.5MG</u>	<u>A212019</u>	<u>001</u>	Dec 12, 2019
<u>AB</u>		<u>1MG</u>	<u>A212019</u>	<u>002</u>	Dec 12, 2019
<u>AB</u>		<u>2MG</u>	<u>A212019</u>	<u>003</u>	Dec 12, 2019
<u>AB</u>	RUBICON	<u>0.5MG</u>	<u>A213942</u>	<u>001</u>	Dec 27, 2024
<u>AB</u>		<u>1MG</u>	<u>A213942</u>	<u>002</u>	Dec 27, 2024
<u>AB</u>		<u>2MG</u>	<u>A213942</u>	<u>003</u>	Dec 27, 2024
<u>AB</u>	SANDOZ	<u>0.5MG</u>	<u>A074700</u>	<u>001</u>	Nov 21, 1996
<u>AB</u>		<u>1MG</u>	<u>A074700</u>	<u>002</u>	Nov 21, 1996
<u>AB</u>	!	<u>2MG</u>	<u>A074700</u>	<u>003</u>	Nov 21, 1996
<u>AB</u>	TARO	<u>0.5MG</u>	<u>A213458</u>	<u>001</u>	Jul 24, 2023
<u>AB</u>		<u>1MG</u>	<u>A213458</u>	<u>002</u>	Jul 24, 2023
<u>AB</u>		<u>2MG</u>	<u>A213458</u>	<u>003</u>	Jul 24, 2023
<u>AB</u>	UPSHER SMITH LABS	<u>0.5MG</u>	<u>A209916</u>	<u>001</u>	Jan 23, 2018
<u>AB</u>		<u>1MG</u>	<u>A209916</u>	<u>002</u>	Jan 23, 2018
<u>AB</u>		<u>2MG</u>	<u>A209916</u>	<u>003</u>	Jan 23, 2018
<u>AB</u>	ZYDUS PHARMS	<u>0.5MG</u>	<u>A202900</u>	<u>001</u>	Apr 30, 2018
<u>AB</u>		<u>1MG</u>	<u>A202900</u>	<u>002</u>	Apr 30, 2018
<u>AB</u>		<u>2MG</u>	<u>A202900</u>	<u>003</u>	Apr 30, 2018
<u>BUMEX</u>					
<u>AB</u>	+ VALIDUS PHARMS	<u>0.5MG</u>	<u>N018225</u>	<u>002</u>	Feb 28, 1983
<u>AB</u>	+	<u>1MG</u>	<u>N018225</u>	<u>001</u>	Feb 28, 1983
<u>AB</u>	+	<u>2MG</u>	<u>N018225</u>	<u>003</u>	Jun 14, 1985

BUPIVACAINE

INJECTABLE, LIPOSOMAL; INJECTION

BUPIVACAINE LIPOSOME

<u>AP</u>	HENGRUI PHARMA	<u>133MG/10ML (13.3MG/ML)</u>	<u>A214348</u>	<u>001</u>	Jul 01, 2024
<u>AP</u>		<u>266MG/20ML (13.3MG/ML)</u>	<u>A214348</u>	<u>002</u>	Jul 01, 2024

EXPAREL

<u>AP</u>	+! PACIRA PHARMS INC	<u>133MG/10ML (13.3MG/ML)</u>	<u>N022496</u>	<u>001</u>	Oct 28, 2011
<u>AP</u>	+!	<u>266MG/20ML (13.3MG/ML)</u>	<u>N022496</u>	<u>002</u>	Oct 28, 2011

BUPIVACAINE HYDROCHLORIDE

INJECTABLE; INJECTION

BUPIVACAINE HYDROCHLORIDE

<u>AP</u>	ASPIRO	<u>0.25%</u>	<u>A217821</u>	<u>001</u>	Mar 27, 2024
<u>AP</u>		<u>0.5%</u>	<u>A217821</u>	<u>002</u>	Mar 27, 2024
<u>AP</u>		<u>0.75%</u>	<u>A217821</u>	<u>003</u>	Mar 27, 2024
<u>AP</u>	EUGIA PHARMA	<u>0.25%</u>	<u>A207183</u>	<u>001</u>	May 13, 2016
<u>AP</u>		<u>0.5%</u>	<u>A207183</u>	<u>002</u>	May 13, 2016
<u>AP</u>	HIKMA PHARMS	<u>0.25%</u>	<u>A205141</u>	<u>001</u>	Feb 11, 2021
<u>AP</u>		<u>0.5%</u>	<u>A205141</u>	<u>002</u>	Feb 11, 2021
<u>AP</u>	HOSPIRA	<u>0.25%</u>	<u>A070583</u>	<u>001</u>	Feb 17, 1987
<u>AP</u>		<u>0.25%</u>	<u>A070590</u>	<u>001</u>	Feb 17, 1987
<u>AP</u>		<u>0.5%</u>	<u>A070584</u>	<u>001</u>	Feb 17, 1986
<u>AP</u>		<u>0.5%</u>	<u>A070597</u>	<u>001</u>	Mar 03, 1987
<u>AP</u>		<u>0.5%</u>	<u>A070609</u>	<u>001</u>	Mar 03, 1987
<u>AP</u>		<u>0.75%</u>	<u>A070585</u>	<u>001</u>	Mar 03, 1987
<u>AP</u>	MEITHEAL	<u>0.25%</u>	<u>A216039</u>	<u>001</u>	Jun 23, 2023
<u>AP</u>		<u>0.25%</u>	<u>A216040</u>	<u>001</u>	Dec 27, 2023
<u>AP</u>		<u>0.5%</u>	<u>A216039</u>	<u>002</u>	Jun 23, 2023
<u>AP</u>		<u>0.5%</u>	<u>A216040</u>	<u>002</u>	Dec 27, 2023
<u>AP</u>		<u>0.75%</u>	<u>A216040</u>	<u>003</u>	Dec 27, 2023
<u>AP</u>	SOMERSET	<u>0.25%</u>	<u>A217792</u>	<u>001</u>	Nov 20, 2023
<u>AP</u>		<u>0.5%</u>	<u>A217792</u>	<u>002</u>	Nov 20, 2023
<u>AP</u>	STERISCIENCE SPECLTS	<u>0.25%</u>	<u>A091503</u>	<u>001</u>	Oct 18, 2011

PRESCRIPTION DRUG PRODUCT LIST

BUPIVACAINE HYDROCHLORIDE

INJECTABLE; INJECTION

BUPIVACAINE HYDROCHLORIDE

<u>AP</u>		<u>0.5%</u>	<u>A091503 002</u>	Oct 18, 2011
	<u>BUPIVACAINE HYDROCHLORIDE PRESERVATIVE FREE</u>			
<u>AP</u>	EUGIA PHARMA	<u>0.25%</u>	<u>A203895 001</u>	Nov 05, 2013
<u>AP</u>		<u>0.5%</u>	<u>A203895 002</u>	Nov 05, 2013
<u>AP</u>		<u>0.75%</u>	<u>A203895 003</u>	Nov 05, 2013
<u>AP</u>	HIKMA PHARMS	<u>0.25%</u>	<u>A204842 001</u>	Feb 11, 2021
<u>AP</u>		<u>0.5%</u>	<u>A204842 002</u>	Feb 11, 2021
<u>AP</u>		<u>0.75%</u>	<u>A204842 003</u>	Feb 11, 2021
<u>AP</u>	STERISCIENCE SPECLTS	<u>0.25%</u>	<u>A091487 002</u>	Oct 18, 2011
<u>AP</u>		<u>0.5%</u>	<u>A091487 001</u>	Oct 18, 2011
<u>AP</u>		<u>0.75%</u>	<u>A091487 003</u>	Oct 18, 2011

MARCAINE HYDROCHLORIDE

<u>AP</u>	+!	HOSPIRA	<u>0.25%</u>	<u>N016964 001</u>	
<u>AP</u>	+!		<u>0.5%</u>	<u>N016964 006</u>	
	<u>MARCAINE HYDROCHLORIDE PRESERVATIVE FREE</u>				
<u>AP</u>	+!	HOSPIRA	<u>0.25%</u>	<u>N016964 012</u>	
<u>AP</u>	+!		<u>0.5%</u>	<u>N016964 005</u>	
<u>AP</u>	+!		<u>0.75%</u>	<u>N016964 009</u>	

SENSORCAINE

<u>AP</u>		FRESENIUS KABI USA	<u>0.25%</u>	<u>A070552 001</u>	May 21, 1986
<u>AP</u>	+		<u>0.25%</u>	<u>N018304 001</u>	
<u>AP</u>			<u>0.5%</u>	<u>A070553 001</u>	May 21, 1986
<u>AP</u>	+		<u>0.5%</u>	<u>N018304 002</u>	
<u>AP</u>			<u>0.75%</u>	<u>A070554 001</u>	May 21, 1986
<u>AP</u>	+		<u>0.75%</u>	<u>N018304 003</u>	

INJECTABLE; SPINAL

BUPIVACAINE HYDROCHLORIDE

<u>AP</u>		B BRAUN MEDICAL INC	<u>0.75%</u>	<u>A209087 001</u>	Apr 02, 2019
<u>AP</u>		HOSPIRA	<u>0.75%</u>	<u>A071810 001</u>	Dec 11, 1987
<u>AP</u>		HUONS	<u>0.75%</u>	<u>A212822 001</u>	Dec 30, 2019

MARCAINE

<u>AP</u>	+!	HOSPIRA	<u>0.75%</u>	<u>N018692 001</u>	May 04, 1984
-----------	----	---------	--------------	--------------------	--------------

BUPIVACAINE HYDROCHLORIDE; EPINEPHRINE

INJECTABLE; INJECTION

BUPIVACAINE HYDROCHLORIDE AND EPINEPHRINE

<u>AP</u>	!	HOSPIRA	<u>0.5%; 0.005MG/ML</u>	<u>A071168 001</u>	Jun 16, 1988
<u>AP</u>			<u>0.5%; 0.005MG/ML</u>	<u>A071170 001</u>	Jun 16, 1988
	!		0.25%; 0.005MG/ML	A071165 001	Jun 16, 1988
			0.25%; 0.005MG/ML	A071167 001	Jun 16, 1988

BUPIVACAINE HYDROCHLORIDE; EPINEPHRINE BITARTRATE

INJECTABLE; INJECTION

MARCAINE HYDROCHLORIDE W/ EPINEPHRINE

<u>AP</u>	+!	HOSPIRA	<u>0.25%; 0.0091MG/ML</u>	<u>N016964 004</u>	
<u>AP</u>	+!		<u>0.5%; 0.0091MG/ML</u>	<u>N016964 008</u>	

MARCAINE HYDROCHLORIDE W/ EPINEPHRINE PRESERVATIVE FREE

<u>AP</u>	+!	HOSPIRA	<u>0.25%; 0.0091MG/ML</u>	<u>N016964 013</u>	
<u>AP</u>	+!		<u>0.5%; 0.0091MG/ML</u>	<u>N016964 007</u>	
<u>AP</u>	+!		<u>0.75%; 0.0091MG/ML</u>	<u>N016964 010</u>	

SENSORCAINE

<u>AP</u>		FRESENIUS KABI USA	<u>0.25%; 0.0091MG/ML</u>	<u>A070966 001</u>	Oct 13, 1987
<u>AP</u>			<u>0.25%; 0.0091MG/ML</u>	<u>A070967 001</u>	Oct 13, 1987
<u>AP</u>			<u>0.5%; 0.0091MG/ML</u>	<u>A070968 001</u>	Oct 13, 1987
<u>AP</u>	+		<u>0.5%; 0.0091MG/ML</u>	<u>N018304 004</u>	Sep 02, 1983
<u>AP</u>	+		<u>0.75%; 0.0091MG/ML</u>	<u>N018304 005</u>	Sep 02, 1983

VIVACAINE

!	SEPTODONT	0.5%; 0.0091MG/ML	A077250 001	Sep 27, 2006
---	-----------	-------------------	-------------	--------------

BUPIVACAINE; MELOXICAM

SOLUTION, EXTENDED RELEASE; PERIARTICULAR

ZYNRELEF KIT

+!	HERON THERAPS INC	200MG/7ML (29.25MG/ML); 6MG/7ML (0.88MG/ML)	N211988 002	May 12, 2021
+!		400MG/14ML (29.25MG/ML); 12MG/14ML (0.88MG/ML)	N211988 004	May 12, 2021

PRESCRIPTION DRUG PRODUCT LIST

BUPRENORPHINE

FILM, EXTENDED RELEASE; TRANSDERMAL

BUPRENORPHINE

<u>AB</u>	ALVOGEN	<u>5MCG/HR</u>	<u>A207490 001</u>	May 17, 2022
<u>AB</u>		<u>10MCG/HR</u>	<u>A207490 002</u>	May 17, 2022
<u>AB</u>		<u>15MCG/HR</u>	<u>A207490 003</u>	May 17, 2022
<u>AB</u>		<u>20MCG/HR</u>	<u>A207490 004</u>	May 17, 2022
<u>AB</u>	AMNEAL	<u>5MCG/HR</u>	<u>A211586 001</u>	Apr 14, 2020
<u>AB</u>		<u>7.5MCG/HR</u>	<u>A211586 002</u>	Apr 14, 2020
<u>AB</u>		<u>10MCG/HR</u>	<u>A211586 003</u>	Apr 14, 2020
<u>AB</u>		<u>15MCG/HR</u>	<u>A211586 004</u>	Apr 14, 2020
<u>AB</u>		<u>20MCG/HR</u>	<u>A211586 005</u>	Apr 14, 2020
<u>AB</u>	DIFGEN PHARMS	<u>5MCG/HR</u>	<u>A210272 001</u>	Sep 23, 2021
<u>AB</u>		<u>7.5MCG/HR</u>	<u>A210272 002</u>	Sep 23, 2021
<u>AB</u>		<u>10MCG/HR</u>	<u>A210272 003</u>	Sep 23, 2021
<u>AB</u>		<u>15MCG/HR</u>	<u>A210272 004</u>	Sep 23, 2021
<u>AB</u>		<u>20MCG/HR</u>	<u>A210272 005</u>	Sep 23, 2021
<u>AB</u>	WATSON LABS TEVA	<u>5MCG/HR</u>	<u>A204937 001</u>	Nov 20, 2018
<u>AB</u>		<u>7.5MCG/HR</u>	<u>A204937 005</u>	Jun 29, 2021
<u>AB</u>		<u>10MCG/HR</u>	<u>A204937 002</u>	Nov 20, 2018
<u>AB</u>		<u>15MCG/HR</u>	<u>A204937 003</u>	Nov 20, 2018
<u>AB</u>		<u>20MCG/HR</u>	<u>A204937 004</u>	Nov 20, 2018

BUTRANS

<u>AB</u>	+	PURDUE PHARMA LP	<u>5MCG/HR</u>	<u>N021306 001</u>	Jun 30, 2010
<u>AB</u>	+		<u>7.5MCG/HR</u>	<u>N021306 005</u>	Jun 30, 2014
<u>AB</u>	+		<u>10MCG/HR</u>	<u>N021306 002</u>	Jun 30, 2010
<u>AB</u>	+		<u>15MCG/HR</u>	<u>N021306 004</u>	Jul 25, 2013
<u>AB</u>	+		<u>20MCG/HR</u>	<u>N021306 003</u>	Jun 30, 2010

SOLUTION, EXTENDED RELEASE; SUBCUTANEOUS

BRIXADI

+	BRAEBURN	8MG/0.16ML (50MG/ML)	N210136 001	May 23, 2023
+		16MG/0.32ML (50MG/ML)	N210136 002	May 23, 2023
+		24MG/0.48ML (50MG/ML)	N210136 003	May 23, 2023
+		32MG/0.64ML (50MG/ML)	N210136 004	May 23, 2023
+		64MG/0.18ML (356MG/ML)	N210136 005	May 23, 2023
+		96MG/0.27ML (356MG/ML)	N210136 006	May 23, 2023
+	!	128MG/0.36ML (356MG/ML)	N210136 007	May 23, 2023

SUBLOCADE

+	INDIVIOR	100MG/0.5ML (100MG/0.5ML)	N209819 001	Nov 30, 2017
+	!	300MG/1.5ML (200MG/ML)	N209819 002	Nov 30, 2017

BUPRENORPHINE HYDROCHLORIDE

FILM; BUCCAL

BELBUCA

+	BDSI	EQ 0.075MG BASE	N207932 001	Oct 23, 2015
+		EQ 0.15MG BASE	N207932 002	Oct 23, 2015
+		EQ 0.3MG BASE	N207932 003	Oct 23, 2015
+		EQ 0.45MG BASE	N207932 004	Oct 23, 2015
+		EQ 0.6MG BASE	N207932 005	Oct 23, 2015
+		EQ 0.75MG BASE	N207932 006	Oct 23, 2015
+	!	EQ 0.9MG BASE	N207932 007	Oct 23, 2015

INJECTABLE; INJECTION

BUPRENORPHINE HYDROCHLORIDE

<u>AP</u>	!	ENDO OPERATIONS	<u>EQ 0.3MG BASE/ML</u>	<u>A206586 001</u>	Jul 28, 2015
<u>AP</u>		HIKMA	<u>EQ 0.3MG BASE/ML</u>	<u>A076931 001</u>	Mar 02, 2005
<u>AP</u>		HOSPIRA	<u>EQ 0.3MG BASE/ML</u>	<u>A074137 001</u>	Jun 03, 1996

TABLET; SUBLINGUAL

BUPRENORPHINE HYDROCHLORIDE

<u>AB</u>		ACTAVIS ELIZABETH	<u>EQ 2MG BASE</u>	<u>A090819 001</u>	Feb 19, 2015
<u>AB</u>			<u>EQ 8MG BASE</u>	<u>A090819 002</u>	Feb 19, 2015
<u>AB</u>		ETHYPHARM	<u>EQ 2MG BASE</u>	<u>A090622 001</u>	Sep 24, 2010
<u>AB</u>			<u>EQ 8MG BASE</u>	<u>A090622 002</u>	Sep 24, 2010
<u>AB</u>		HIKMA	<u>EQ 2MG BASE</u>	<u>A078633 001</u>	Oct 08, 2009
<u>AB</u>	!		<u>EQ 8MG BASE</u>	<u>A078633 002</u>	Oct 08, 2009
<u>AB</u>		RHODES PHARMS	<u>EQ 2MG BASE</u>	<u>A207276 001</u>	Mar 27, 2017
<u>AB</u>			<u>EQ 8MG BASE</u>	<u>A207276 002</u>	Mar 27, 2017
<u>AB</u>		RUBICON	<u>EQ 2MG BASE</u>	<u>A090279 001</u>	Jun 10, 2015
<u>AB</u>			<u>EQ 8MG BASE</u>	<u>A090279 002</u>	Jun 10, 2015
<u>AB</u>		SUN PHARM	<u>EQ 2MG BASE</u>	<u>A201760 001</u>	Jan 29, 2016
<u>AB</u>			<u>EQ 8MG BASE</u>	<u>A201760 002</u>	Jan 29, 2016

PRESCRIPTION DRUG PRODUCT LIST

BUPRENORPHINE HYDROCHLORIDE; NALOXONE HYDROCHLORIDE

FILM;BUCCAL, SUBLINGUAL

BUPRENORPHINE HYDROCHLORIDE AND NALOXONE HYDROCHLORIDE

<u>AB</u>	ALVOGEN	<u>EQ 2MG BASE;EQ 0.5MG BASE</u>	<u>A205954 001</u>	Jan 24, 2019
<u>AB</u>		<u>EQ 4MG BASE;EQ 1MG BASE</u>	<u>A205954 002</u>	Jan 24, 2019
<u>AB</u>		<u>EQ 8MG BASE;EQ 2MG BASE</u>	<u>A205954 003</u>	Jan 24, 2019
<u>AB</u>		<u>EQ 12MG BASE;EQ 3MG BASE</u>	<u>A205954 004</u>	Jan 24, 2019
<u>AB</u>	DIFGEN PHARMS	<u>EQ 8MG BASE;EQ 2MG BASE</u>	<u>A212756 001</u>	Jun 02, 2022
<u>AB</u>		<u>EQ 12MG BASE;EQ 3MG BASE</u>	<u>A212756 002</u>	Jun 02, 2022
<u>AB</u>	DR REDDYS LABS SA	<u>EQ 2MG BASE;EQ 0.5MG BASE</u>	<u>A205299 001</u>	Jun 14, 2018
<u>AB</u>		<u>EQ 4MG BASE;EQ 1MG BASE</u>	<u>A205806 001</u>	Jun 14, 2018
<u>AB</u>		<u>EQ 8MG BASE;EQ 2MG BASE</u>	<u>A205299 002</u>	Jun 14, 2018
<u>AB</u>		<u>EQ 12MG BASE;EQ 3MG BASE</u>	<u>A205806 002</u>	Jun 14, 2018
<u>AB</u>	MYLAN TECHNOLOGIES	<u>EQ 2MG BASE;EQ 0.5MG BASE</u>	<u>A211785 001</u>	Apr 17, 2020
<u>AB</u>		<u>EQ 4MG BASE;EQ 1MG BASE</u>	<u>A211785 002</u>	Apr 17, 2020
<u>AB</u>		<u>EQ 8MG BASE;EQ 2MG BASE</u>	<u>A207607 001</u>	Jun 14, 2018
<u>AB</u>		<u>EQ 12MG BASE;EQ 3MG BASE</u>	<u>A207607 002</u>	Jun 14, 2018

SUBOXONE

<u>AB</u>	+ INDIVIOR	<u>EQ 2MG BASE;EQ 0.5MG BASE</u>	<u>N022410 001</u>	Aug 30, 2010
<u>AB</u>	+	<u>EQ 4MG BASE;EQ 1MG BASE</u>	<u>N022410 003</u>	Aug 10, 2012
<u>AB</u>	+	<u>EQ 8MG BASE;EQ 2MG BASE</u>	<u>N022410 002</u>	Aug 30, 2010
<u>AB</u>	+!	<u>EQ 12MG BASE;EQ 3MG BASE</u>	<u>N022410 004</u>	Aug 10, 2012

TABLET;SUBLINGUAL

BUPRENORPHINE HYDROCHLORIDE AND NALOXONE HYDROCHLORIDE

<u>AB</u>	ACTAVIS ELIZABETH	<u>EQ 2MG BASE;EQ 0.5MG BASE</u>	<u>A091422 001</u>	Feb 22, 2013
<u>AB</u>		<u>EQ 8MG BASE;EQ 2MG BASE</u>	<u>A091422 002</u>	Feb 22, 2013
<u>AB</u>	ALKEM LABS LTD	<u>EQ 2MG BASE;EQ 0.5MG BASE</u>	<u>A214930 001</u>	Jun 15, 2021
<u>AB</u>		<u>EQ 8MG BASE;EQ 2MG BASE</u>	<u>A214930 002</u>	Jun 15, 2021
<u>AB</u>	AMNEAL PHARMS	<u>EQ 2MG BASE;EQ 0.5MG BASE</u>	<u>A203136 001</u>	Feb 22, 2013
<u>AB</u>		<u>EQ 8MG BASE;EQ 2MG BASE</u>	<u>A203136 002</u>	Feb 22, 2013
<u>AB</u>	ETHYPHARM USA CORP	<u>EQ 2MG BASE;EQ 0.5MG BASE</u>	<u>A204431 001</u>	Oct 16, 2015
<u>AB</u>		<u>EQ 8MG BASE;EQ 2MG BASE</u>	<u>A204431 002</u>	Oct 16, 2015
<u>AB</u>	HIKMA	<u>EQ 2MG BASE;EQ 0.5MG BASE</u>	<u>A203326 001</u>	Jun 27, 2014
<u>AB</u>	!	<u>EQ 8MG BASE;EQ 2MG BASE</u>	<u>A203326 002</u>	Jun 27, 2014
<u>AB</u>	LANNETT CO INC	<u>EQ 2MG BASE;EQ 0.5MG BASE</u>	<u>A205022 001</u>	Sep 19, 2016
<u>AB</u>		<u>EQ 8MG BASE;EQ 2MG BASE</u>	<u>A205022 002</u>	Sep 19, 2016
<u>AB</u>	RHODES PHARMS	<u>EQ 2MG BASE;EQ 0.5MG BASE</u>	<u>A205601 001</u>	Mar 30, 2020
<u>AB</u>		<u>EQ 8MG BASE;EQ 2MG BASE</u>	<u>A205601 002</u>	Mar 30, 2020
<u>AB</u>	SPECGX LLC	<u>EQ 2MG BASE;EQ 0.5MG BASE</u>	<u>A207000 001</u>	Dec 13, 2017
<u>AB</u>		<u>EQ 8MG BASE;EQ 2MG BASE</u>	<u>A207000 002</u>	Dec 13, 2017
<u>AB</u>	SUN PHARM	<u>EQ 2MG BASE;EQ 0.5MG BASE</u>	<u>A201633 001</u>	Aug 05, 2016
<u>AB</u>		<u>EQ 8MG BASE;EQ 2MG BASE</u>	<u>A201633 002</u>	Aug 05, 2016
<u>AB</u>	WES PHARMA INC	<u>EQ 2MG BASE;EQ 0.5MG BASE</u>	<u>A209069 001</u>	Jul 17, 2020
<u>AB</u>		<u>EQ 8MG BASE;EQ 2MG BASE</u>	<u>A209069 002</u>	Jul 17, 2020

ZUBSOLV

+	OREXO US INC	EQ 0.7MG BASE;EQ 0.18MG BASE	N204242 006	Oct 04, 2016
+		EQ 1.4MG BASE;EQ 0.36MG BASE	N204242 001	Jul 03, 2013
+		EQ 2.9MG BASE;EQ 0.71MG BASE	N204242 005	Jun 04, 2015
+		EQ 5.7MG BASE;EQ 1.4MG BASE	N204242 002	Jul 03, 2013
+		EQ 8.6MG BASE;EQ 2.1MG BASE	N204242 003	Dec 11, 2014
+	!	EQ 11.4MG BASE;EQ 2.9MG BASE	N204242 004	Dec 11, 2014

BUPROPION HYDROBROMIDE

TABLET, EXTENDED RELEASE;ORAL

APLENZIN

+	BAUSCH	174MG	N022108 001	Apr 23, 2008
+		348MG	N022108 002	Apr 23, 2008
+	!	522MG	N022108 003	Apr 23, 2008

BUPROPION HYDROCHLORIDE

TABLET;ORAL

BUPROPION HYDROCHLORIDE

<u>AB</u>	ALEMbic	<u>75MG</u>	<u>A203013 001</u>	Jun 08, 2018
<u>AB</u>		<u>100MG</u>	<u>A203013 002</u>	Jun 08, 2018
<u>AB</u>	APNAR PHARMA LP	<u>75MG</u>	<u>A075584 001</u>	Feb 07, 2000
<u>AB</u>		<u>100MG</u>	<u>A075584 002</u>	Feb 07, 2000
<u>AB</u>	APOTEX INC	<u>75MG</u>	<u>A076143 001</u>	Jan 17, 2006
<u>AB</u>	!	<u>100MG</u>	<u>A076143 002</u>	Jan 17, 2006
<u>AB</u>	AUROBINDO PHARMA USA	<u>75MG</u>	<u>A075491 001</u>	Apr 17, 2000
<u>AB</u>		<u>100MG</u>	<u>A075491 002</u>	Apr 17, 2000
<u>AB</u>	CADILA PHARMS LTD	<u>75MG</u>	<u>A208606 001</u>	Jan 16, 2020
<u>AB</u>		<u>100MG</u>	<u>A208606 002</u>	Jan 16, 2020
<u>AB</u>	HERITAGE PHARMA	<u>75MG</u>	<u>A206975 001</u>	Aug 19, 2016

PRESCRIPTION DRUG PRODUCT LIST

BUPROPION HYDROCHLORIDE

TABLET;ORAL

BUPROPION HYDROCHLORIDE

<u>AB</u>		<u>100MG</u>	<u>A206975</u>	<u>002</u>	Aug 19, 2016
<u>AB</u>	MICRO LABS	<u>75MG</u>	<u>A207403</u>	<u>001</u>	Apr 17, 2020
<u>AB</u>		<u>100MG</u>	<u>A207403</u>	<u>002</u>	Apr 17, 2020

TABLET, EXTENDED RELEASE;ORAL

BUPROPION HYDROCHLORIDE

<u>AB1</u>	ACTAVIS LABS FL INC	<u>100MG</u>	<u>A079095</u>	<u>001</u>	Mar 24, 2009
<u>AB1</u>		<u>150MG</u>	<u>A079095</u>	<u>002</u>	Mar 24, 2009
<u>AB1</u>		<u>200MG</u>	<u>A079095</u>	<u>003</u>	Mar 24, 2009
<u>AB1</u>	ANNORA PHARMA	<u>100MG</u>	<u>A216800</u>	<u>001</u>	May 31, 2023
<u>AB1</u>		<u>150MG</u>	<u>A216800</u>	<u>002</u>	May 31, 2023
<u>AB1</u>		<u>200MG</u>	<u>A216800</u>	<u>003</u>	May 31, 2023
<u>AB1</u>	EPIC PHARMA LLC	<u>100MG</u>	<u>A075932</u>	<u>001</u>	Nov 25, 2003
<u>AB1</u>		<u>150MG</u>	<u>A075932</u>	<u>002</u>	Mar 22, 2004
<u>AB1</u>		<u>200MG</u>	<u>A075932</u>	<u>003</u>	Jun 22, 2005
<u>AB1</u>	GRANULES	<u>100MG</u>	<u>A218385</u>	<u>001</u>	Oct 17, 2024
<u>AB1</u>		<u>150MG</u>	<u>A218385</u>	<u>002</u>	Oct 17, 2024
<u>AB1</u>		<u>200MG</u>	<u>A218385</u>	<u>003</u>	Oct 17, 2024
<u>AB1</u>	IMPAX LABS	<u>100MG</u>	<u>A075913</u>	<u>001</u>	Jan 28, 2004
<u>AB1</u>		<u>150MG</u>	<u>A075913</u>	<u>002</u>	Mar 22, 2004
<u>AB1</u>	PRINSTON INC	<u>100MG</u>	<u>A202304</u>	<u>001</u>	May 26, 2015
<u>AB1</u>		<u>150MG</u>	<u>A202304</u>	<u>002</u>	May 26, 2015
<u>AB1</u>		<u>200MG</u>	<u>A202304</u>	<u>003</u>	May 26, 2015
<u>AB1</u>	SCIEGEN PHARMS INC	<u>100MG</u>	<u>A205794</u>	<u>001</u>	Mar 01, 2016
<u>AB1</u>		<u>150MG</u>	<u>A205794</u>	<u>002</u>	Mar 01, 2016
<u>AB1</u>		<u>200MG</u>	<u>A205794</u>	<u>003</u>	Mar 01, 2016
<u>AB1</u>	YICHANG HUMANWELL	<u>100MG</u>	<u>A211347</u>	<u>001</u>	Oct 16, 2018
<u>AB1</u>		<u>150MG</u>	<u>A211347</u>	<u>002</u>	Oct 16, 2018
<u>AB1</u>		<u>200MG</u>	<u>A211347</u>	<u>003</u>	Oct 16, 2018

WELLBUTRIN SR

<u>AB1</u>	+	GLAXOSMITHKLINE	<u>100MG</u>	<u>N020358</u>	<u>002</u>	Oct 04, 1996
<u>AB1</u>	+		<u>150MG</u>	<u>N020358</u>	<u>003</u>	Oct 04, 1996
<u>AB1</u>	+	!	<u>200MG</u>	<u>N020358</u>	<u>004</u>	Jun 14, 2002

BUPROPION HYDROCHLORIDE

<u>AB2</u>	!	ACTAVIS LABS FL INC	<u>150MG</u>	<u>A079094</u>	<u>001</u>	Mar 24, 2009
<u>AB2</u>		IMPAX LABS	<u>150MG</u>	<u>A075914</u>	<u>001</u>	May 27, 2004
<u>AB2</u>		SANDOZ	<u>150MG</u>	<u>A077475</u>	<u>001</u>	Mar 12, 2008
<u>AB2</u>		SCIEGEN PHARMS INC	<u>150MG</u>	<u>A206122</u>	<u>001</u>	Aug 17, 2016
<u>AB2</u>		YICHANG HUMANWELL	<u>150MG</u>	<u>A216766</u>	<u>001</u>	Jan 09, 2023
<u>AB3</u>		ACCORD HLTHCARE	<u>150MG</u>	<u>A210497</u>	<u>001</u>	Oct 31, 2018
<u>AB3</u>			<u>300MG</u>	<u>A210497</u>	<u>002</u>	Oct 31, 2018
<u>AB3</u>		ACTAVIS LABS FL INC	<u>150MG</u>	<u>A077715</u>	<u>001</u>	Nov 26, 2008
<u>AB3</u>		ANBISON LAB	<u>150MG</u>	<u>A207224</u>	<u>001</u>	Jun 30, 2017
<u>AB3</u>			<u>300MG</u>	<u>A207224</u>	<u>002</u>	Jun 30, 2017
<u>AB3</u>		GRANULES	<u>150MG</u>	<u>A215568</u>	<u>001</u>	Feb 02, 2022
<u>AB3</u>			<u>300MG</u>	<u>A215568</u>	<u>002</u>	Feb 02, 2022
<u>AB3</u>		GRAVITI PHARMS	<u>150MG</u>	<u>A211020</u>	<u>001</u>	Jan 28, 2019
<u>AB3</u>			<u>300MG</u>	<u>A211020</u>	<u>002</u>	Jan 28, 2019
<u>AB3</u>		LUPIN LTD	<u>150MG</u>	<u>A090693</u>	<u>001</u>	Apr 06, 2017
<u>AB3</u>			<u>300MG</u>	<u>A090693</u>	<u>002</u>	Apr 06, 2017
<u>AB3</u>		SCIEGEN PHARMS INC	<u>150MG</u>	<u>A207479</u>	<u>001</u>	Apr 12, 2017
<u>AB3</u>			<u>300MG</u>	<u>A207479</u>	<u>002</u>	Apr 12, 2017
<u>AB3</u>		SINOTHERAPEUTICS INC	<u>150MG</u>	<u>A208652</u>	<u>001</u>	Aug 21, 2017
<u>AB3</u>			<u>300MG</u>	<u>A208652</u>	<u>002</u>	Aug 21, 2017
<u>AB3</u>		SUN PHARM	<u>150MG</u>	<u>A200216</u>	<u>001</u>	Nov 30, 2020
<u>AB3</u>			<u>300MG</u>	<u>A203650</u>	<u>001</u>	Dec 31, 2020
<u>AB3</u>		TWI PHARMS	<u>150MG</u>	<u>A210081</u>	<u>001</u>	Nov 03, 2017
<u>AB3</u>			<u>300MG</u>	<u>A210081</u>	<u>002</u>	Nov 03, 2017
<u>AB3</u>		WATSON LABS INC	<u>150MG</u>	<u>A077285</u>	<u>001</u>	Nov 26, 2008
<u>AB3</u>			<u>300MG</u>	<u>A077285</u>	<u>002</u>	Aug 15, 2008
<u>AB3</u>		WOCKHARDT LTD	<u>150MG</u>	<u>A202189</u>	<u>001</u>	Nov 21, 2012
<u>AB3</u>			<u>300MG</u>	<u>A202189</u>	<u>002</u>	Jan 28, 2022
<u>AB3</u>		YICHANG HUMANWELL	<u>150MG</u>	<u>A210015</u>	<u>001</u>	Jun 14, 2018
<u>AB3</u>			<u>300MG</u>	<u>A210015</u>	<u>002</u>	Jun 14, 2018
<u>AB3</u>		ZHEJIANG JUTAI PHARM	<u>150MG</u>	<u>A211200</u>	<u>002</u>	Apr 29, 2020
<u>AB3</u>			<u>300MG</u>	<u>A211200</u>	<u>001</u>	Sep 05, 2019
<u>AB3</u>		ZYDUS PHARMS	<u>150MG</u>	<u>A201567</u>	<u>002</u>	Jul 23, 2018
<u>AB3</u>			<u>300MG</u>	<u>A201567</u>	<u>001</u>	Jan 17, 2014

PRESCRIPTION DRUG PRODUCT LIST

BUPROPION HYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL

WELLBUTRIN XL

AB3	+	BAUSCH	150MG	N021515	001	Aug 28, 2003
AB3	+	!	300MG	N021515	002	Aug 28, 2003
		FORFIVO XL				
	+	TWI PHARMS	450MG	N022497	001	Nov 10, 2011

BUPROPION HYDROCHLORIDE; DEXTROMETHORPHAN HYDROBROMIDE

TABLET, EXTENDED RELEASE;ORAL

AUVELITY

+	AXSOME	105MG;45MG	N215430	001	Aug 18, 2022
---	--------	------------	---------	-----	--------------

BUPROPION HYDROCHLORIDE; NALTREXONE HYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL

CONTRAVE

+	NALPROPION	90MG;8MG	N200063	001	Sep 10, 2014
---	------------	----------	---------	-----	--------------

BUSPIRONE HYDROCHLORIDE

TABLET;ORAL

BUSPIRONE HYDROCHLORIDE

AB		ACCORD HLTHCARE	5MG	A202557	001	Dec 30, 2014
AB			7.5MG	A202557	002	Dec 30, 2014
AB			10MG	A202557	003	Dec 30, 2014
AB	!		15MG	A202557	004	Dec 30, 2014
AB			30MG	A202557	005	Dec 30, 2014
AB		AIPING PHARM INC	5MG	A202087	001	Dec 16, 2015
AB			10MG	A202087	002	Dec 16, 2015
AB			15MG	A202087	003	Dec 16, 2015
AB			30MG	A202087	004	Dec 16, 2015
AB		AUROBINDO PHARMA LTD	5MG	A078246	001	Feb 27, 2009
AB			7.5MG	A078246	005	Feb 21, 2020
AB			10MG	A078246	002	Feb 27, 2009
AB			15MG	A078246	003	Feb 27, 2009
AB			30MG	A078246	004	Feb 27, 2009
AB		EPIC PHARMA LLC	5MG	A208972	001	Apr 16, 2019
AB			7.5MG	A208972	002	Apr 16, 2019
AB			10MG	A208972	003	Apr 16, 2019
AB			15MG	A208972	004	Apr 16, 2019
AB			30MG	A208972	005	Apr 16, 2019
AB		HERITAGE PHARMA	5MG	A204582	001	Sep 18, 2015
AB			10MG	A204582	002	Sep 18, 2015
AB			15MG	A204582	003	Sep 18, 2015
AB			30MG	A204582	004	Sep 18, 2015
AB		IMPAX LABS INC	5MG	A074253	001	Mar 28, 2001
AB			10MG	A074253	002	Mar 28, 2001
AB			15MG	A074253	003	Mar 13, 2002
AB		INVENTIA HLTHCARE	5MG	A209696	001	May 03, 2018
AB			7.5MG	A209696	002	May 03, 2018
AB			10MG	A209696	003	May 03, 2018
AB			15MG	A209696	004	May 03, 2018
AB			30MG	A209696	005	May 03, 2018
AB		MYLAN	5MG	A076008	003	Mar 01, 2002
AB			7.5MG	A076008	002	Jul 08, 2013
AB			10MG	A076008	004	Mar 01, 2002
AB			15MG	A076008	005	Mar 28, 2001
AB			30MG	A076008	001	Jun 28, 2001
AB		OXFORD PHARMS	5MG	A075388	001	May 09, 2002
AB			10MG	A075388	002	May 09, 2002
AB			15MG	A075388	003	May 09, 2002
AB			30MG	A078302	001	Dec 17, 2007
AB		RUBICON	5MG	A075521	001	Apr 05, 2002
AB			7.5MG	A075521	004	Mar 16, 2021
AB			10MG	A075521	002	Apr 05, 2002
AB			15MG	A075521	003	Apr 05, 2002
AB			30MG	A075521	005	Mar 16, 2021
AB		STRIDES PHARMA	5MG	A202330	001	Aug 25, 2014
AB			7.5MG	A202330	005	Feb 17, 2017
AB			10MG	A202330	002	Aug 25, 2014
AB			15MG	A202330	003	Aug 25, 2014
AB			30MG	A202330	004	Aug 25, 2014
AB		TEVA	5MG	A075022	001	Feb 28, 2002
AB			10MG	A075022	002	Feb 28, 2002

PRESCRIPTION DRUG PRODUCT LIST

BUSPIRONE HYDROCHLORIDE

TABLET; ORAL

BUSPIRONE HYDROCHLORIDE

<u>AB</u>		<u>15MG</u>	<u>A075022 003</u>	Feb 28, 2002
<u>AB</u>		<u>30MG</u>	<u>A075022 004</u>	Mar 25, 2004
<u>AB</u>	UNICHEM	<u>5MG</u>	<u>A210907 001</u>	Nov 14, 2019
<u>AB</u>		<u>10MG</u>	<u>A210907 002</u>	Nov 14, 2019
<u>AB</u>		<u>15MG</u>	<u>A210907 003</u>	Nov 14, 2019
<u>AB</u>		<u>30MG</u>	<u>A210907 004</u>	Nov 14, 2019
<u>AB</u>	ZYDUS PHARMS	<u>5MG</u>	<u>A078888 001</u>	Feb 07, 2014
<u>AB</u>		<u>7.5MG</u>	<u>A078888 005</u>	Mar 21, 2023
<u>AB</u>		<u>10MG</u>	<u>A078888 002</u>	Feb 07, 2014
<u>AB</u>		<u>15MG</u>	<u>A078888 003</u>	Feb 07, 2014
<u>AB</u>		<u>30MG</u>	<u>A078888 004</u>	Feb 07, 2014

BUSULFAN

INJECTABLE; INJECTION

BUSULFAN

<u>AP</u>	ACCORD HLTHCARE INC	<u>6MG/ML</u>	<u>A210148 001</u>	Feb 22, 2019
<u>AP</u>	AMNEAL	<u>6MG/ML</u>	<u>A209580 001</u>	Dec 18, 2017
<u>AP</u>	APOTEX	<u>6MG/ML</u>	<u>A210448 001</u>	May 07, 2019
<u>AP</u>	HOSPIRA	<u>6MG/ML</u>	<u>A205672 001</u>	Jul 31, 2018
<u>AP</u>	MEITHEAL	<u>6MG/ML</u>	<u>A212127 001</u>	Oct 23, 2020
<u>AP</u>	MYLAN INSTITUTIONAL	<u>6MG/ML</u>	<u>A208536 001</u>	Nov 20, 2017
<u>AP</u>	PHARMSCIENCE INC	<u>6MG/ML</u>	<u>A207050 001</u>	Mar 24, 2017
<u>AP</u>	PRINSTON INC	<u>6MG/ML</u>	<u>A215235 001</u>	Sep 11, 2024
<u>AP</u>	SHILPA	<u>6MG/ML</u>	<u>A210931 001</u>	Apr 18, 2019

BUSULFEX

<u>AP</u>	+! OTSUKA PHARM	<u>6MG/ML</u>	<u>N020954 001</u>	Feb 04, 1999
	TABLET; ORAL			
	MYLERAN			
	+! WAYLIS THERAP	2MG	N009386 001	

BUTOCONAZOLE NITRATE

CREAM; VAGINAL

GYNAZOLE-1

!	PADAGIS ISRAEL	2%	A200923 001	May 18, 2012
---	----------------	----	-------------	--------------

BUTORPHANOL TARTRATE

INJECTABLE; INJECTION

BUTORPHANOL TARTRATE

<u>AP</u>	HIKMA	<u>2MG/ML</u>	<u>A075046 001</u>	Aug 12, 1998
<u>AP</u>	HIKMA FARMACEUTICA	<u>1MG/ML</u>	<u>A078400 001</u>	May 01, 2009
<u>AP</u>		<u>2MG/ML</u>	<u>A078400 002</u>	May 01, 2009

BUTORPHANOL TARTRATE PRESERVATIVE FREE

<u>AP</u>	HIKMA	<u>1MG/ML</u>	<u>A075045 001</u>	Aug 12, 1998
<u>AP</u>		<u>2MG/ML</u>	<u>A075045 002</u>	Aug 12, 1998
<u>AP</u>	! HOSPIRA	<u>1MG/ML</u>	<u>A074626 001</u>	Jan 23, 1997
<u>AP</u>	!	<u>2MG/ML</u>	<u>A074626 002</u>	Jan 23, 1997

SPRAY, METERED; NASAL

BUTORPHANOL TARTRATE

<u>AB</u>	APOTEX	<u>1MG/SPRAY</u>	<u>A075499 001</u>	Dec 04, 2002
<u>AB</u>	HIKMA	<u>1MG/SPRAY</u>	<u>A075824 001</u>	Mar 12, 2002
<u>AB</u>	! RISING	<u>1MG/SPRAY</u>	<u>A075759 001</u>	Aug 08, 2001

CABAZITAXEL

SOLUTION; INTRAVENOUS

CABAZITAXEL

<u>AP</u>	ACCORD HLTHCARE	<u>60MG/1.5ML (40MG/ML)</u>	<u>A207693 001</u>	Oct 26, 2022
<u>AP</u>	DR REDDYS	<u>60MG/1.5ML (40MG/ML)</u>	<u>A207718 001</u>	Feb 10, 2023

JEVTANA KIT

<u>AP</u>	+! SANOFI AVENTIS US	<u>60MG/1.5ML (40MG/ML)</u>	<u>N201023 001</u>	Jun 17, 2010
-----------	----------------------	-----------------------------	--------------------	--------------

CABERGOLINE

TABLET; ORAL

CABERGOLINE

<u>AB</u>	! INGENUS PHARMS LLC	<u>0.5MG</u>	<u>A204735 001</u>	Aug 01, 2018
<u>AB</u>	IVAX SUB TEVA PHARMS	<u>0.5MG</u>	<u>A077750 001</u>	Mar 07, 2007
<u>AB</u>	SKG PHARMA	<u>0.5MG</u>	<u>A218618 001</u>	Jun 26, 2024
<u>AB</u>	SOMERSET THERAPS LLC	<u>0.5MG</u>	<u>A218109 001</u>	Sep 09, 2024
<u>AB</u>	STRIDES PHARMA	<u>0.5MG</u>	<u>A076310 001</u>	Dec 29, 2005

PRESCRIPTION DRUG PRODUCT LIST

CABOTEGRAVIR

SUSPENSION, EXTENDED RELEASE; INTRAMUSCULAR

APRETUDE

+! VIIV HLTHCARE 600MG/3ML (200MG/ML) N215499 001 Dec 20, 2021

CABOTEGRAVIR SODIUM

TABLET; ORAL

VOCABRIA

+! VIIV HLTHCARE EQ 30MG BASE N212887 001 Jan 21, 2021

CABOTEGRAVIR; RILPIVIRINE

SUSPENSION, EXTENDED RELEASE; INTRAMUSCULAR

CABENUVA KIT

+! VIIV HLTHCARE 400MG/2ML (200MG/ML); 600MG/2ML (300MG/ML) N212888 001 Jan 21, 2021

+! 600MG/3ML (200MG/ML); 900MG/3ML (300MG/ML) N212888 002 Jan 21, 2021

CABOZANTINIB S-MALATE

CAPSULE; ORAL

COMETRIQ

+ EXELIXIS EQ 20MG BASE N203756 001 Nov 29, 2012

+! EQ 80MG BASE N203756 002 Nov 29, 2012

TABLET; ORAL

CABOMETYX

+ EXELIXIS INC EQ 20MG BASE N208692 001 Apr 25, 2016

+ EQ 40MG BASE N208692 002 Apr 25, 2016

+! EQ 60MG BASE N208692 003 Apr 25, 2016

CAFFEINE CITRATE

SOLUTION; INTRAVENOUS

CAFCITAP +! HIKMA EQ 30MG BASE/3ML (EQ 10MG BASE/ML) N020793 001 Sep 21, 1999CAFFEINE CITRATEAP AM REGENT EQ 30MG BASE/3ML (EQ 10MG BASE/ML) A077906 001 May 15, 2007AP EUGIA PHARMA EQ 30MG BASE/3ML (EQ 10MG BASE/ML) A205013 001 Sep 22, 2015AP EXELA PHARMA EQ 30MG BASE/3ML (EQ 10MG BASE/ML) A077233 001 Sep 21, 2006

SCIENCE

AP FRESENIUS KABI USA EQ 30MG BASE/3ML (EQ 10MG BASE/ML) A077997 001 Jul 20, 2007AP MICRO LABS EQ 30MG BASE/3ML (EQ 10MG BASE/ML) A207400 001 Dec 14, 2017AP SAGENT PHARMS EQ 30MG BASE/3ML (EQ 10MG BASE/ML) A090827 001 Aug 29, 2012

SOLUTION; ORAL

CAFFEINE CITRATEAA ! EXELA PHARMA EQ 30MG BASE/3ML (EQ 10MG BASE/ML) A077304 001 Sep 21, 2006AA FRESENIUS KABI USA EQ 30MG BASE/3ML (EQ 10MG BASE/ML) A078002 001 Jan 31, 2008AA MICRO LABS EQ 30MG BASE/3ML (EQ 10MG BASE/ML) A213202 001 Dec 16, 2019AA SAGENT PHARMS EQ 30MG BASE/3ML (EQ 10MG BASE/ML) A091102 001 Aug 29, 2012AA SUN PHARM EQ 30MG BASE/3ML (EQ 10MG BASE/ML) A090357 001 Sep 30, 2009CAFFEINE; ERGOTAMINE TARTRATE

SUPPOSITORY; RECTAL

MIGERGOT

+! COSETTE 100MG; 2MG A086557 001 Oct 04, 1983

TABLET; ORAL

ERGOTAMINE TARTRATE AND CAFFEINE

! MIKART 100MG; 1MG A040590 001 Sep 16, 2005

CALCIFEDIOL

CAPSULE, EXTENDED RELEASE; ORAL

RAYALDEE

+! EIRGEN 0.03MG N208010 001 Jun 17, 2016

CALCIPOTRIENE

AEROSOL, FOAM; TOPICAL

SORILUX

+! MAYNE PHARMA 0.005% N022563 001 Oct 06, 2010

CREAM; TOPICAL

CALCIPOTRIENEAB GLENMARK SPECLT 0.005% A205772 001 Jun 09, 2015DOVONEXAB +! LEO PHARMA AS 0.005% N020554 001 Jul 22, 1996

OINTMENT; TOPICAL

CALCIPOTRIENEAB ! GLENMARK PHARMS INC 0.005% A090633 001 Mar 24, 2010DOVONEXAB + LEO PHARMA AS 0.005% N020273 001 Dec 29, 1993

PRESCRIPTION DRUG PRODUCT LIST

CALCIPOTRIENE

SOLUTION; TOPICAL

CALCIPOTRIENE

AT	CHARTWELL RX	<u>0.005%</u>	<u>A207163</u>	<u>001</u>	Dec 26, 2017	
AT	!	COSETTE	<u>0.005%</u>	<u>A078468</u>	<u>001</u>	Mar 24, 2011
AT	FOUGERA PHARMS	<u>0.005%</u>	<u>A078305</u>	<u>001</u>	May 06, 2008	

CALCITONIN SALMON

INJECTABLE; INJECTION

CALCITONIN-SALMON

AP	CIPLA	<u>200 IU/ML</u>	<u>A213766</u>	<u>001</u>	Sep 20, 2024
AP	CUSTOPHARM INC	<u>200 IU/ML</u>	<u>A212416</u>	<u>001</u>	May 14, 2021
AP	DR REDDYS	<u>200 IU/ML</u>	<u>A215715</u>	<u>001</u>	Apr 11, 2024
AP	ENDO OPERATIONS	<u>200 IU/ML</u>	<u>A209358</u>	<u>001</u>	Nov 10, 2021
AP	FRESENIUS KABI USA	<u>200 IU/ML</u>	<u>A212675</u>	<u>001</u>	Dec 05, 2024

MIACALCIN

AP	+	MYLAN IRELAND LTD	<u>200 IU/ML</u>	<u>N017808</u>	<u>002</u>	Mar 29, 1991
-----------	---	-------------------	------------------	----------------	------------	--------------

SPRAY, METERED; NASAL

CALCITONIN-SALMON

AB	!	APOTEX INC	<u>200 IU/SPRAY</u>	<u>A076396</u>	<u>001</u>	Nov 17, 2008
AB		ENDO OPERATIONS	<u>200 IU/SPRAY</u>	<u>A076979</u>	<u>001</u>	Jun 08, 2009

CALCITRIOL

CAPSULE; ORAL

CALCITRIOL

AB		AMNEAL PHARMS	<u>0.25MCG</u>	<u>A203289</u>	<u>002</u>	Jun 14, 2017
AB			<u>0.5MCG</u>	<u>A203289</u>	<u>001</u>	Jun 14, 2017
AB		BIONPHARMA	<u>0.25MCG</u>	<u>A091174</u>	<u>001</u>	May 24, 2013
AB			<u>0.5MCG</u>	<u>A091174</u>	<u>002</u>	May 24, 2013
AB		HIKMA	<u>0.25MCG</u>	<u>A076917</u>	<u>001</u>	Mar 27, 2006
AB		STRIDES SOFTGELS	<u>0.25MCG</u>	<u>A091356</u>	<u>001</u>	Dec 12, 2014
AB	!		<u>0.5MCG</u>	<u>A091356</u>	<u>002</u>	Dec 12, 2014
AB		TEVA	<u>0.25MCG</u>	<u>A075765</u>	<u>001</u>	Oct 12, 2001
AB			<u>0.5MCG</u>	<u>A075765</u>	<u>002</u>	Oct 12, 2001

ROCALTROL

AB	+	ESJAY PHARMA	<u>0.25MCG</u>	<u>N018044</u>	<u>001</u>	
AB	+		<u>0.5MCG</u>	<u>N018044</u>	<u>002</u>	

INJECTABLE; INJECTION

CALCITRIOL

AP		GLAND PHARMA LTD	<u>0.001MG/ML</u>	<u>A211030</u>	<u>001</u>	Feb 03, 2020
AP	!	LONG GROVE PHARMS	<u>0.001MG/ML</u>	<u>A078066</u>	<u>001</u>	Jan 29, 2008

OINTMENT; TOPICAL

VECTICAL

	+	GALDERMA LABS LP	3MCG/GM	N022087	001	Jan 23, 2009
--	---	------------------	---------	---------	-----	--------------

SOLUTION; ORAL

CALCITRIOL

AA		ANDA REPOSITORY	<u>1MCG/ML</u>	<u>A203973</u>	<u>001</u>	Jul 28, 2023
AA		HIKMA	<u>1MCG/ML</u>	<u>A076242</u>	<u>001</u>	Jul 18, 2003
AA		RISING	<u>1MCG/ML</u>	<u>A209798</u>	<u>001</u>	Nov 21, 2018

ROCALTROL

AA	+	ESJAY PHARMA	<u>1MCG/ML</u>	<u>N021068</u>	<u>001</u>	Nov 20, 1998
-----------	---	--------------	----------------	----------------	------------	--------------

CALCIUM ACETATE

CAPSULE; ORAL

CALCIUM ACETATE

AB		CHARTWELL RX	<u>667MG</u>	<u>A091312</u>	<u>001</u>	Jun 01, 2012
AB		HERITAGE PHARMS INC	<u>667MG</u>	<u>A202315</u>	<u>001</u>	Jun 29, 2015
AB		HIKMA	<u>667MG</u>	<u>A077728</u>	<u>001</u>	Feb 26, 2008
AB	!	INVAGEN PHARMS	<u>667MG</u>	<u>A203135</u>	<u>001</u>	Feb 07, 2013
AB		LUPIN LTD	<u>667MG</u>	<u>A202127</u>	<u>001</u>	Jul 09, 2015
AB		NOSTRUM LABS INC	<u>667MG</u>	<u>A203179</u>	<u>001</u>	Oct 26, 2015
AB		SQUARE PHARMS	<u>667MG</u>	<u>A217205</u>	<u>001</u>	Mar 13, 2023
AB		SUVEN PHARMS	<u>667MG</u>	<u>A211038</u>	<u>001</u>	Feb 21, 2020

TABLET; ORAL

CALCIUM ACETATE

AB	!	CHARTWELL MOLECULAR	<u>667MG</u>	<u>A202420</u>	<u>001</u>	Feb 05, 2013
AB		HERITAGE PHARMS INC	<u>667MG</u>	<u>A202885</u>	<u>001</u>	Jan 22, 2015
AB		PADAGIS US	<u>667MG</u>	<u>A091561</u>	<u>001</u>	Apr 13, 2011

PRESCRIPTION DRUG PRODUCT LIST

CALCIUM CHLORIDE

INJECTABLE; INJECTION

CALCIUM CHLORIDE 10%

AP	AM REGENT	<u>100MG/ML</u>	<u>A209088 001</u>	Jul 27, 2017
AP	AMNEAL	<u>100MG/ML</u>	<u>A217524 001</u>	Jul 08, 2024
AP	EXTROVIS	<u>100MG/ML</u>	<u>A219342 001</u>	Nov 27, 2024
AP	INTL MEDICATION SYS	<u>100MG/ML</u>	<u>A203477 001</u>	May 09, 2018
AP	MEDEFIL INC	<u>100MG/ML</u>	<u>A211553 001</u>	May 01, 2019
AP	SOMERSET	<u>100MG/ML</u>	<u>A218133 001</u>	Dec 17, 2024

CALCIUM CHLORIDE 10% IN PLASTIC CONTAINER

AP	+ ! HOSPIRA	<u>100MG/ML</u>	<u>N021117 001</u>	Jan 28, 2000
-----------	--------------------	-----------------	--------------------	--------------

CALCIUM CHLORIDE; DEXTROSE; GLUTATHIONE DISULFIDE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE; SODIUM PHOSPHATE

SOLUTION; IRRIGATION

BSS PLUS

+ !	ALCON	0.154MG/ML; 0.92MG/ML; 0.184MG/ML; 0.2MG/M L; 0.38MG/ML; 2.1MG/ML; 7.14MG/ML; 0.42MG/M L	N018469 001	
------------	-------	--	-------------	--

CALCIUM CHLORIDE; DEXTROSE; LACTIC ACID; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE

INJECTABLE; INJECTION

PRISMASOL B22GK 4/0 IN PLASTIC CONTAINER

+ !	BAXTER HLTHCARE CORP	N/A/1000ML; 20GM/1000ML; 5.4GM/1000ML; 3.0 5GM/1000ML; 0.314GM/1000ML; 2.21GM/1000ML ; 7.07GM/1000ML (5000ML)	N021703 011	Oct 10, 2008
------------	-------------------------	---	-------------	--------------

PRISMASOL BGK 0/2.5 IN PLASTIC CONTAINER

+ !	BAXTER HLTHCARE CORP	3.68GM/1000ML; 20GM/1000ML; 5.4GM/1000ML; 3.05GM/1000ML; N/A/1000ML; 3.09GM/1000ML; 6.46GM/1000ML (5000ML)	N021703 006	Oct 25, 2006
------------	-------------------------	--	-------------	--------------

PRISMASOL BGK 2/0 IN PLASTIC CONTAINER

+ !	BAXTER HLTHCARE CORP	N/A/1000ML; 20GM/1000ML; 5.4GM/1000ML; 2.0 3GM/1000ML; 0.157GM/1000ML; 3.09GM/1000ML ; 6.46GM/1000ML (5000ML)	N021703 002	Oct 25, 2006
------------	-------------------------	---	-------------	--------------

PRISMASOL BGK 2/3.5 IN PLASTIC CONTAINER

+ !	BAXTER HLTHCARE CORP	5.15GM/1000ML; 20GM/1000ML; 5.4GM/1000ML; 2.03GM/1000ML; 0.157GM/1000ML; 3.09GM/100 0ML; 6.46GM/1000ML (5000ML)	N021703 003	Oct 25, 2006
------------	-------------------------	---	-------------	--------------

PRISMASOL BGK 4/0/1.2 IN PLASTIC CONTAINER

+ !	BAXTER HLTHCARE CORP	N/A/1000ML; 20GM/1000ML; 5.4GM/1000ML; 2.4 4GM/1000ML; 0.314GM/1000ML; 3.09GM/1000ML ; 6.46GM/1000ML (5000ML)	N021703 015	Oct 10, 2008
------------	-------------------------	---	-------------	--------------

PRISMASOL BGK 4/2.5 IN PLASTIC CONTAINER

+ !	BAXTER HLTHCARE CORP	3.68GM/1000ML; 20GM/1000ML; 5.4GM/1000ML; 3.05GM/1000ML; 0.314GM/1000ML; 3.09GM/100 0ML; 6.46GM/1000ML (5000ML)	N021703 004	Oct 25, 2006
------------	-------------------------	---	-------------	--------------

PRISMASOL BK 0/0/1.2 IN PLASTIC CONTAINER

+ !	BAXTER HLTHCARE CORP	N/A/1000ML; N/A/1000ML; 5.4GM/1000ML; 2.44 GM/1000ML; N/A/1000ML; 3.09GM/1000ML; 6.46 GM/1000ML (5000ML)	N021703 014	Oct 10, 2008
------------	-------------------------	--	-------------	--------------

CALCIUM CHLORIDE; DEXTROSE; MAGNESIUM CHLORIDE; SODIUM CHLORIDE; SODIUM LACTATE

SOLUTION; INTRAPERITONEAL

DELFLX W/ DEXTROSE 1.5% IN PLASTIC CONTAINER

AT	+ ! FRESENIUS MEDCL	<u>25.7MG/100ML; 1.5GM/100ML; 15.2MG/100ML; 5 67MG/100ML; 392MG/100ML</u>	<u>N018883 001</u>	Nov 30, 1984
-----------	----------------------------	---	--------------------	--------------

DELFLX W/ DEXTROSE 1.5% LOW MAGNESIUM IN PLASTIC CONTAINER

AT	+ ! FRESENIUS MEDCL	<u>25.7MG/100ML; 1.5GM/100ML; 5.08MG/100ML; 5 38MG/100ML; 448MG/100ML</u>	<u>N018883 004</u>	Nov 30, 1984
-----------	----------------------------	---	--------------------	--------------

DELFLX W/ DEXTROSE 1.5% LOW MAGNESIUM LOW CALCIUM IN PLASTIC CONTAINER

AT	+ ! FRESENIUS MEDCL	<u>18.4MG/100ML; 1.5GM/100ML; 5.08MG/100ML; 5 38MG/100ML; 448MG/100ML</u>	<u>N020171 001</u>	Aug 19, 1992
-----------	----------------------------	---	--------------------	--------------

DELFLX W/ DEXTROSE 2.5% IN PLASTIC CONTAINER

AT	+ ! FRESENIUS MEDCL	<u>25.7MG/100ML; 2.5GM/100ML; 15.2MG/100ML; 5 67MG/100ML; 392MG/100ML</u>	<u>N018883 002</u>	Nov 30, 1984
-----------	----------------------------	---	--------------------	--------------

DELFLX W/ DEXTROSE 2.5% LOW MAGNESIUM IN PLASTIC CONTAINER

AT	+ ! FRESENIUS MEDCL	<u>25.7MG/100ML; 2.5GM/100ML; 5.08MG/100ML; 5 38MG/100ML; 448MG/100ML</u>	<u>N018883 005</u>	Nov 30, 1984
-----------	----------------------------	---	--------------------	--------------

DELFLX W/ DEXTROSE 2.5% LOW MAGNESIUM LOW CALCIUM IN PLASTIC CONTAINER

AT	+ ! FRESENIUS MEDCL	<u>18.4MG/100ML; 2.5GM/100ML; 5.08MG/100ML; 5 38MG/100ML; 448MG/100ML</u>	<u>N020171 002</u>	Aug 19, 1992
-----------	----------------------------	---	--------------------	--------------

DELFLX W/ DEXTROSE 4.25% IN PLASTIC CONTAINER

AT	+ ! FRESENIUS MEDCL	<u>25.7MG/100ML; 4.25GM/100ML; 15.2MG/100ML; 567MG/100ML; 392MG/100ML</u>	<u>N018883 003</u>	Nov 30, 1984
-----------	----------------------------	---	--------------------	--------------

DELFLX W/ DEXTROSE 4.25% LOW MAGNESIUM IN PLASTIC CONTAINER

AT	+ ! FRESENIUS MEDCL	<u>25.7MG/100ML; 4.25GM/100ML; 5.08MG/100ML; 538MG/100ML; 448MG/100ML</u>	<u>N018883 006</u>	Nov 30, 1984
-----------	----------------------------	---	--------------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

CALCIUM CHLORIDE; DEXTROSE; MAGNESIUM CHLORIDE; SODIUM CHLORIDE; SODIUM LACTATE
SOLUTION; INTRAPERITONEAL

<u>DELFLX W/ DEXTROSE 4.25% LOW MAGNESIUM LOW CALCIUM IN PLASTIC CONTAINER</u>					
<u>AT</u>	<u>+!</u>	<u>FRESENIUS MEDCL</u>	<u>18.4MG/100ML;4.25GM/100ML;5.08MG/100ML;</u> <u>538MG/100ML;448MG/100ML</u>	<u>N020171 003</u>	<u>Aug 19, 1992</u>
<u>DIANEAL LOW CALCIUM W/ DEXTROSE 1.5% IN PLASTIC CONTAINER</u>					
<u>AT</u>	<u>+</u>	<u>BAXTER HLTHCARE</u>	<u>18.3MG/100ML;1.5GM/100ML;5.08MG/100ML;5</u> <u>38MG/100ML;448MG/100ML</u>	<u>N020183 001</u>	<u>Dec 04, 1992</u>
<u>DIANEAL LOW CALCIUM W/ DEXTROSE 2.5% IN PLASTIC CONTAINER</u>					
<u>AT</u>	<u>+</u>	<u>BAXTER HLTHCARE</u>	<u>18.3MG/100ML;2.5GM/100ML;5.08MG/100ML;5</u> <u>38MG/100ML;448MG/100ML</u>	<u>N020183 002</u>	<u>Dec 04, 1992</u>
<u>DIANEAL LOW CALCIUM W/ DEXTROSE 4.25% IN PLASTIC CONTAINER</u>					
<u>AT</u>	<u>+</u>	<u>BAXTER HLTHCARE</u>	<u>18.3MG/100ML;4.25GM/100ML;5.08MG/100ML;</u> <u>538MG/100ML;448MG/100ML</u>	<u>N020183 004</u>	<u>Dec 04, 1992</u>
<u>DIANEAL LOW CALCIUM W/ DEXTROSE 1.5% IN PLASTIC CONTAINER</u>					
<u>AT</u>	<u>+</u>	<u>BAXTER HLTHCARE</u>	<u>18.3MG/100ML;1.5GM/100ML;5.08MG/100ML;5</u> <u>38MG/100ML;448MG/100ML</u>	<u>N017512 012</u>	<u>Jan 10, 1989</u>
<u>DIANEAL LOW CALCIUM W/ DEXTROSE 2.5% IN PLASTIC CONTAINER</u>					
<u>AT</u>	<u>+</u>	<u>BAXTER HLTHCARE</u>	<u>18.3MG/100ML;2.5GM/100ML;5.08MG/100ML;5</u> <u>38MG/100ML;448MG/100ML</u>	<u>N017512 013</u>	<u>Jul 11, 1990</u>
<u>DIANEAL LOW CALCIUM W/ DEXTROSE 3.5% IN PLASTIC CONTAINER</u>					
<u>AT</u>	<u>+</u>	<u>BAXTER HLTHCARE</u>	<u>18.3MG/100ML;3.5GM/100ML;5.08MG/100ML;5</u> <u>38MG/100ML;448MG/100ML</u>	<u>N017512 014</u>	<u>Jul 11, 1990</u>
<u>DIANEAL LOW CALCIUM W/ DEXTROSE 4.25% IN PLASTIC CONTAINER</u>					
<u>AT</u>	<u>+</u>	<u>BAXTER HLTHCARE</u>	<u>18.3MG/100ML;4.25GM/100ML;5.08MG/100ML;</u> <u>538MG/100ML;448MG/100ML</u>	<u>N017512 015</u>	<u>Jul 11, 1990</u>
<u>DIANEAL PD-2 W/ DEXTROSE 1.5% IN PLASTIC CONTAINER</u>					
<u>AT</u>	<u>+</u>	<u>BAXTER HLTHCARE</u>	<u>25.7MG/100ML;1.5GM/100ML;5.08MG/100ML;5</u> <u>38MG/100ML;448MG/100ML</u>	<u>N017512 004</u>	
<u>AT</u>	<u>+</u>		<u>25.7MG/100ML;1.5GM/100ML;5.08MG/100ML;5</u> <u>38MG/100ML;448MG/100ML</u>	<u>N020163 001</u>	<u>Dec 04, 1992</u>
<u>DIANEAL PD-2 W/ DEXTROSE 2.5% IN PLASTIC CONTAINER</u>					
<u>AT</u>	<u>+</u>	<u>BAXTER HLTHCARE</u>	<u>25.7MG/100ML;2.5GM/100ML;5.08MG/100ML;5</u> <u>38MG/100ML;448MG/100ML</u>	<u>N017512 005</u>	
<u>AT</u>	<u>+</u>		<u>25.7MG/100ML;2.5GM/100ML;5.08MG/100ML;5</u> <u>38MG/100ML;448MG/100ML</u>	<u>N020163 002</u>	<u>Dec 04, 1992</u>
<u>DIANEAL PD-2 W/ DEXTROSE 4.25% IN PLASTIC CONTAINER</u>					
<u>AT</u>	<u>+</u>	<u>BAXTER HLTHCARE</u>	<u>25.7MG/100ML;4.25GM/100ML;5.08MG/100ML;</u> <u>538MG/100ML;448MG/100ML</u>	<u>N017512 006</u>	
<u>AT</u>	<u>+</u>		<u>25.7MG/100ML;4.25GM/100ML;5.08MG/100ML;</u> <u>538MG/100ML;448MG/100ML</u>	<u>N020163 003</u>	<u>Dec 04, 1992</u>

CALCIUM CHLORIDE; DEXTROSE; MAGNESIUM SULFATE; POTASSIUM CHLORIDE; SODIUM BICARBONATE; SODIUM
CHLORIDE; SODIUM PHOSPHATE, DIBASIC, HEPTAHYDRATE

INJECTABLE; INTRATHECAL

ELLIOTTS B SOLUTION

<u>+!</u>	<u>LUKARE MEDICAL LLC</u>	<u>0.2MG/ML;0.8MG/ML;0.3MG/ML;0.3MG/ML;1.9</u> <u>MG/ML;7.3MG/ML;0.2MG/ML</u>	<u>N020577 001</u>	<u>Sep 27, 1996</u>
-----------	---------------------------	--	--------------------	---------------------

CALCIUM CHLORIDE; DEXTROSE; POTASSIUM CHLORIDE; SODIUM CHLORIDE; SODIUM LACTATE

INJECTABLE; INJECTION

<u>DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER</u>					
<u>AP</u>	<u>+</u>	<u>ICU MEDICAL INC</u>	<u>20MG/100ML;5GM/100ML;30MG/100ML;600MG/1</u> <u>00ML;310MG/100ML</u>	<u>N017608 001</u>	
<u>DEXTROSE 5% IN LACTATED RINGER'S IN PLASTIC CONTAINER</u>					
<u>AP</u>		<u>B BRAUN</u>	<u>20MG/100ML;5GM/100ML;30MG/100ML;600MG/1</u> <u>00ML;310MG/100ML</u>	<u>N019634 003</u>	<u>Feb 24, 1988</u>
<u>LACTATED RINGER'S AND DEXTROSE 5% IN PLASTIC CONTAINER</u>					
<u>AP</u>		<u>BAXTER HLTHCARE</u>	<u>20MG/100ML;5GM/100ML;30MG/100ML;600MG/1</u> <u>00ML;310MG/100ML</u>	<u>N016679 001</u>	
<u>POTASSIUM CHLORIDE 15MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER</u>					
<u>AP</u>		<u>BAXTER HLTHCARE</u>	<u>20MG/100ML;5GM/100ML;254MG/100ML;600MG/</u> <u>100ML;310MG/100ML</u>	<u>N019367 006</u>	<u>Apr 05, 1985</u>
<u>POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER</u>					
<u>AP</u>		<u>BAXTER HLTHCARE</u>	<u>20MG/100ML;5GM/100ML;179MG/100ML;600MG/</u> <u>100ML;310MG/100ML</u>	<u>N019367 004</u>	<u>Apr 05, 1985</u>
<u>AP</u>			<u>20MG/100ML;5GM/100ML;328MG/100ML;600MG/</u> <u>100ML;310MG/100ML</u>	<u>N019367 005</u>	<u>Apr 05, 1985</u>
<u>AP</u>	<u>+</u>	<u>ICU MEDICAL INC</u>	<u>20MG/100ML;5GM/100ML;179MG/100ML;600MG/</u> <u>100ML;310MG/100ML</u>	<u>N019685 002</u>	<u>Oct 17, 1988</u>
<u>POTASSIUM CHLORIDE 30MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER</u>					
<u>AP</u>		<u>BAXTER HLTHCARE</u>	<u>20MG/100ML;5GM/100ML;254MG/100ML;600MG/</u> <u>100ML;310MG/100ML</u>	<u>N019367 007</u>	<u>Apr 05, 1985</u>
<u>POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER</u>					
<u>AP</u>		<u>BAXTER HLTHCARE</u>	<u>20MG/100ML;5GM/100ML;328MG/100ML;600MG/</u> <u>100ML;310MG/100ML</u>	<u>N019367 008</u>	<u>Apr 05, 1985</u>
<u>DEXTROSE 2.5% IN HALF-STRENGTH LACTATED RINGER'S IN PLASTIC CONTAINER</u>					
		<u>B BRAUN</u>	<u>10MG/100ML;2.5GM/100ML;15MG/100ML;</u>	<u>N019634 001</u>	<u>Feb 24, 1988</u>

PRESCRIPTION DRUG PRODUCT LIST

CALCIUM CHLORIDE; DEXTROSE; POTASSIUM CHLORIDE; SODIUM CHLORIDE; SODIUM LACTATE

INJECTABLE; INJECTION

DEXTROSE 2.5% IN HALF-STRENGTH LACTATED RINGER'S IN PLASTIC CONTAINER

300MG/100ML; 160MG/100ML

POTASSIUM CHLORIDE 10MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER

BAXTER HLTHCARE

20MG/100ML; 5GM/100ML; 105MG/100ML; 600MG/
100ML; 310MG/100ML

N019367 002 Apr 05, 1985

20MG/100ML; 5GM/100ML; 179MG/100ML; 600MG/
100ML; 310MG/100ML

N019367 003 Apr 05, 1985

POTASSIUM CHLORIDE 5MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER

BAXTER HLTHCARE

20MG/100ML; 5GM/100ML; 105MG/100ML; 600MG/
100ML; 310MG/100ML

N019367 001 Apr 05, 1985

CALCIUM CHLORIDE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE

INJECTABLE; INJECTION

TPN ELECTROLYTES IN PLASTIC CONTAINER

+! HOSPIRA

16.5MG/ML; 25.4MG/ML; 74.6MG/ML; 121MG/ML;
16.1MG/ML

N018895 001 Jul 20, 1984

CALCIUM CHLORIDE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE; SODIUM CITRATE

SOLUTION; IRRIGATION

BALANCED SALT

AT

B BRAUN

0.48MG/ML; 0.3MG/ML; 0.75MG/ML; 3.9MG/ML; 6
.4MG/ML; 1.7MG/MLA091387 001 Feb 03, 2010BSS

AT

+!

ALCON

0.48MG/ML; 0.3MG/ML; 0.75MG/ML; 3.9MG/ML; 6
.4MG/ML; 1.7MG/MLN020742 001 Dec 10, 1997CALCIUM CHLORIDE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE; SODIUM PHOSPHATE

INJECTABLE; INJECTION

PHOXILLUM B22K 4/0 IN PLASTIC CONTAINER

+! BAXTER HLTHCARE
CORPN/A/1000ML; 3.05GM/1000ML; 0.314GM/1000ML
; 2.21GM/1000ML; 6.95GM/1000ML; 0.187GM/10
00ML (5000ML)

N207026 002 Jan 13, 2015

PHOXILLUM BK 4/2.5 IN PLASTIC CONTAINER

+! BAXTER HLTHCARE
CORP3.68GM/1000ML; 3.05GM/1000ML; 0.314GM/100
0ML

N207026 001 Jan 13, 2015

; 3.09GM/1000ML; 6.34GM/1000ML; 0.187GM/10

CALCIUM CHLORIDE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE

SOLUTION; PERFUSION, CARDIAC

CARDIOPLEGIC IN PLASTIC CONTAINER

AT

BAXTER HLTHCARE

17.6MG/100ML; 325.3MG/100ML; 119.3MG/100M
L: 643MG/100MLA075323 001 Apr 21, 2000

AT

FRESENIUS KABI USA

17.6MG/100ML; 325.3MG/100ML; 119.3MG/100M
L: 643MG/100MLA214623 001 Feb 18, 2022PLEGISOL IN PLASTIC CONTAINER

AT

+!

HOSPIRA

17.6MG/100ML; 325.3MG/100ML; 119.3MG/100M
L: 643MG/100MLN018608 001 Feb 26, 1982CALCIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM CHLORIDE

INJECTABLE; INJECTION

RINGER'S IN PLASTIC CONTAINER

AP

B BRAUN

33MG/100ML; 30MG/100ML; 860MG/100MLN020002 001 Apr 17, 1992

AP

BAXTER HLTHCARE

33MG/100ML; 30MG/100ML; 860MG/100MLN016693 001

AP

ICU MEDICAL INC

33MG/100ML; 30MG/100ML; 860MG/100MLN018251 001

SOLUTION; IRRIGATION

RINGER'S IN PLASTIC CONTAINER

AT

B BRAUN

33MG/100ML; 30MG/100ML; 860MG/100MLN018156 001

AT

BAXTER HLTHCARE

33MG/100ML; 30MG/100ML; 860MG/100MLN018495 001 Feb 19, 1982

AT

ICU MEDICAL INC

33MG/100ML; 30MG/100ML; 860MG/100MLN017635 001CALCIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM CHLORIDE; SODIUM LACTATE

INJECTABLE; INJECTION

LACTATED RINGER'S IN PLASTIC CONTAINER

AP

B BRAUN

20MG/100ML; 30MG/100ML; 600MG/100ML; 310MG
/100MLN019632 001 Feb 29, 1988

AP

+!

BAXTER HLTHCARE

20MG/100ML; 30MG/100ML; 600MG/100ML; 310MG
/100MLN016682 001

AP

FRESENIUS KABI USA

20MG/100ML; 30MG/100ML; 600MG/100ML; 310MG
/100MLA209338 001 Jan 28, 2019

AP

ICU MEDICAL INC

20MG/100ML; 30MG/100ML; 600MG/100ML; 310MG
/100MLN017641 001

SOLUTION; IRRIGATION

LACTATED RINGER'S IN PLASTIC CONTAINER

AT

+!

B BRAUN

20MG/100ML; 30MG/100ML; 600MG/100ML; 310MG
/100MLN018681 001 Dec 27, 1982

PRESCRIPTION DRUG PRODUCT LIST

CALCIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM CHLORIDE; SODIUM LACTATE

SOLUTION; IRRIGATION

LACTATED RINGER'S IN PLASTIC CONTAINER

<u>AT</u>	BAXTER HLTHCARE	<u>20MG/100ML; 30MG/100ML; 600MG/100ML; 310MG/100ML</u>	<u>N018494 001</u>	Feb 19, 1982
<u>AT</u>	+!	<u>20MG/100ML; 30MG/100ML; 600MG/100ML; 310MG/100ML</u>	<u>N018921 001</u>	Apr 03, 1984
<u>AT</u>	+! ICU MEDICAL INC	<u>20MG/100ML; 30MG/100ML; 600MG/100ML; 310MG/100ML</u>	<u>N019416 001</u>	Jan 17, 1986

CALCIUM GLUCONATE

SOLUTION; INTRAVENOUS

CALCIUM GLUCONATE

<u>AP</u>	AMNEAL	<u>1GM/10ML (100MG/ML)</u>	<u>A216611 001</u>	May 10, 2024
<u>AP</u>		<u>5GM/50ML (100MG/ML)</u>	<u>A216611 002</u>	May 10, 2024
<u>AP</u>		<u>10GM/100ML (100MG/ML)</u>	<u>A216611 003</u>	May 10, 2024
<u>AP</u>	B BRAUN MEDICAL INC	<u>10GM/100ML (100MG/ML)</u>	<u>A216541 001</u>	Aug 21, 2023
<u>AP</u>	+! FRESENIUS KABI USA	<u>1GM/10ML (100MG/ML)</u>	<u>N208418 001</u>	Jun 15, 2017
<u>AP</u>	+!	<u>5GM/50ML (100MG/ML)</u>	<u>N208418 002</u>	Jun 15, 2017
<u>AP</u>	+!	<u>10GM/100ML (100MG/ML)</u>	<u>N208418 003</u>	Jun 15, 2017
<u>AP</u>	NIVAGEN PHARMS INC	<u>1GM/10ML (100MG/ML)</u>	<u>A213071 001</u>	Oct 14, 2022
<u>AP</u>		<u>5GM/50ML (100MG/ML)</u>	<u>A213071 002</u>	Oct 19, 2023
<u>AP</u>		<u>10GM/100ML (100MG/ML)</u>	<u>A213071 003</u>	Apr 30, 2024
<u>AP</u>	SOMERSET	<u>5GM/50ML (100MG/ML)</u>	<u>A217689 001</u>	Oct 19, 2023
<u>AP</u>		<u>10GM/100ML (100MG/ML)</u>	<u>A217689 002</u>	Oct 19, 2023

CALCIUM GLUCONATE IN SODIUM CHLORIDE

<u>AP</u>	AMNEAL	<u>1GM/50ML (20MG/ML)</u>	<u>A217174 001</u>	Sep 05, 2023
<u>AP</u>		<u>2GM/100ML (20MG/ML)</u>	<u>A217174 002</u>	Sep 05, 2023
<u>AP</u>	+! FRESENIUS KABI USA	<u>1GM/50ML (20MG/ML)</u>	<u>N208418 004</u>	Jun 17, 2021
<u>AP</u>	+!	<u>2GM/100ML (20MG/ML)</u>	<u>N208418 005</u>	Jun 17, 2021
	+! HQ SPCLT PHARMA	1GM/50ML (20MG/ML)	N210906 001	Oct 29, 2018
	+!	1GM/100ML (10MG/ML)	N210906 003	Jun 04, 2021
	+!	2GM/100ML (20MG/ML)	N210906 002	Oct 29, 2018

CALCIUM OXYBATE; MAGNESIUM OXYBATE; POTASSIUM OXYBATE; SODIUM OXYBATE

SOLUTION; ORAL

XYWAV

	+! JAZZ	0.234GM/ML; 0.096GM/ML; 0.13GM/ML; 0.04GM/ML	N212690 001	Jul 21, 2020
--	---------	--	-------------	--------------

CANAGLIFLOZIN

TABLET; ORAL

INVOKANA

	+ JANSSEN PHARMS	100MG	N204042 001	Mar 29, 2013
	+!	300MG	N204042 002	Mar 29, 2013

CANAGLIFLOZIN; METFORMIN HYDROCHLORIDE

TABLET; ORAL

INVOKAMET

	+ JANSSEN PHARMS	50MG; 500MG	N204353 001	Aug 08, 2014
	+	50MG; 1GM	N204353 002	Aug 08, 2014
	+	150MG; 500MG	N204353 003	Aug 08, 2014
	+!	150MG; 1GM	N204353 004	Aug 08, 2014

TABLET, EXTENDED RELEASE; ORAL

INVOKAMET XR

	+ JANSSEN PHARMS	50MG; 500MG	N205879 001	Sep 20, 2016
	+	50MG; 1GM	N205879 002	Sep 20, 2016
	+	150MG; 500MG	N205879 003	Sep 20, 2016
	+!	150MG; 1GM	N205879 004	Sep 20, 2016

CANDESARTAN CILEXETIL

TABLET; ORAL

ATACAND

<u>AB</u>	+ ANI PHARMS	<u>4MG</u>	<u>N020838 001</u>	Jun 04, 1998
<u>AB</u>	+	<u>8MG</u>	<u>N020838 002</u>	Jun 04, 1998
<u>AB</u>	+	<u>16MG</u>	<u>N020838 003</u>	Jun 04, 1998
<u>AB</u>	+!	<u>32MG</u>	<u>N020838 004</u>	Jun 04, 1998

CANDESARTAN CILEXETIL

<u>AB</u>	ALEMBIC	<u>4MG</u>	<u>A210302 001</u>	Dec 04, 2018
<u>AB</u>		<u>8MG</u>	<u>A210302 002</u>	Dec 04, 2018
<u>AB</u>		<u>16MG</u>	<u>A210302 003</u>	Dec 04, 2018
<u>AB</u>		<u>32MG</u>	<u>A209119 001</u>	Jun 20, 2017
<u>AB</u>	MACLEODS PHARMS LTD	<u>4MG</u>	<u>A203813 001</u>	Dec 05, 2016
<u>AB</u>		<u>8MG</u>	<u>A203813 002</u>	Dec 05, 2016
<u>AB</u>		<u>16MG</u>	<u>A203813 003</u>	Dec 05, 2016
<u>AB</u>		<u>32MG</u>	<u>A203813 004</u>	Dec 05, 2016

PRESCRIPTION DRUG PRODUCT LIST

CANDESARTAN CILEXETIL

TABLET; ORAL

CANDESARTAN CILEXETIL

<u>AB</u>	MYLAN	<u>4MG</u>	<u>A078702 001</u>	May 03, 2013
<u>AB</u>		<u>8MG</u>	<u>A078702 002</u>	May 03, 2013
<u>AB</u>		<u>16MG</u>	<u>A078702 003</u>	May 03, 2013
<u>AB</u>		<u>32MG</u>	<u>A078702 004</u>	May 03, 2013
<u>AB</u>	PRINSTON INC	<u>8MG</u>	<u>A206233 001</u>	Aug 21, 2023
<u>AB</u>		<u>16MG</u>	<u>A206233 002</u>	Aug 21, 2023
<u>AB</u>		<u>32MG</u>	<u>A206233 003</u>	Aug 21, 2023

CANDESARTAN CILEXETIL; HYDROCHLOROTHIAZIDE

TABLET; ORAL

ATACAND HCT

<u>AB</u>	+	ANI PHARMS	<u>16MG;12.5MG</u>	<u>N021093 001</u>	Sep 05, 2000
<u>AB</u>	+		<u>32MG;12.5MG</u>	<u>N021093 002</u>	Sep 05, 2000
<u>AB</u>	+	!	<u>32MG;25MG</u>	<u>N021093 003</u>	May 16, 2008

CANDESARTAN CILEXETIL AND HYDROCHLOROTHIAZIDE

<u>AB</u>		DR REDDYS LABS LTD	<u>16MG;12.5MG</u>	<u>A202965 001</u>	Jun 03, 2013
<u>AB</u>			<u>32MG;12.5MG</u>	<u>A202965 002</u>	Jun 03, 2013
<u>AB</u>			<u>32MG;25MG</u>	<u>A202965 003</u>	Jun 03, 2013
<u>AB</u>		MACLEODS PHARMS LTD	<u>16MG;12.5MG</u>	<u>A204100 001</u>	Feb 27, 2015
<u>AB</u>			<u>32MG;12.5MG</u>	<u>A204100 002</u>	Feb 27, 2015
<u>AB</u>			<u>32MG;25MG</u>	<u>A204100 003</u>	Feb 27, 2015
<u>AB</u>		MYLAN	<u>16MG;12.5MG</u>	<u>A090704 001</u>	Dec 04, 2012
<u>AB</u>			<u>32MG;12.5MG</u>	<u>A090704 002</u>	Dec 04, 2012
<u>AB</u>			<u>32MG;25MG</u>	<u>A090704 003</u>	Dec 04, 2012
<u>AB</u>		PRINSTON INC	<u>16MG;12.5MG</u>	<u>A207455 001</u>	Apr 11, 2018
<u>AB</u>			<u>32MG;12.5MG</u>	<u>A207455 002</u>	Apr 11, 2018
<u>AB</u>			<u>32MG;25MG</u>	<u>A207455 003</u>	Apr 11, 2018
<u>AB</u>		ZYDUS LIFESCIENCES	<u>16MG;12.5MG</u>	<u>A203466 001</u>	Nov 27, 2017
<u>AB</u>			<u>32MG;12.5MG</u>	<u>A203466 002</u>	Nov 27, 2017
<u>AB</u>			<u>32MG;25MG</u>	<u>A203466 003</u>	Nov 27, 2017

CANGRELOR

POWDER; INTRAVENOUS

KENGREAL

+	!	CHIESI	50MG/VIAL	N204958 001	Jun 22, 2015
---	---	--------	-----------	-------------	--------------

CANNABIDIOL

SOLUTION; ORAL

EPIDIOLEX

+	!	JAZZ PHARMS RES	100MG/ML	N210365 001	Sep 28, 2018
---	---	-----------------	----------	-------------	--------------

CANTHARIDIN

SOLUTION; TOPICAL

YCANTH

+	!	VERRICA PHARMS	0.7%	N212905 001	Jul 21, 2023
---	---	----------------	------	-------------	--------------

CAPECITABINE

TABLET; ORAL

CAPECITABINE

<u>AB</u>		ACCORD HLTHCARE	<u>150MG</u>	<u>A202593 001</u>	Apr 23, 2015
<u>AB</u>			<u>500MG</u>	<u>A202593 002</u>	Apr 23, 2015
<u>AB</u>		ALKEM LABS LTD	<u>150MG</u>	<u>A207652 001</u>	Nov 24, 2017
<u>AB</u>			<u>500MG</u>	<u>A207652 002</u>	Nov 24, 2017
<u>AB</u>		DR REDDYS	<u>150MG</u>	<u>A204345 001</u>	Dec 04, 2020
<u>AB</u>			<u>500MG</u>	<u>A204345 002</u>	Dec 04, 2020
<u>AB</u>		EUGIA PHARMA	<u>150MG</u>	<u>A210604 001</u>	Apr 17, 2018
<u>AB</u>			<u>500MG</u>	<u>A210604 002</u>	Apr 17, 2018
<u>AB</u>		HETERO LABS LTD V	<u>150MG</u>	<u>A210203 001</u>	Mar 05, 2024
<u>AB</u>			<u>500MG</u>	<u>A210203 002</u>	Mar 05, 2024
<u>AB</u>		MSN	<u>150MG</u>	<u>A209365 001</u>	Jul 02, 2018
<u>AB</u>			<u>500MG</u>	<u>A209365 002</u>	Jul 02, 2018
<u>AB</u>		RELIANCE LIFE	<u>150MG</u>	<u>A211724 001</u>	Apr 27, 2020
<u>AB</u>			<u>500MG</u>	<u>A211724 002</u>	Apr 27, 2020
<u>AB</u>		RISING	<u>150MG</u>	<u>A090943 001</u>	Aug 08, 2014
<u>AB</u>			<u>500MG</u>	<u>A090943 002</u>	Aug 08, 2014
<u>AB</u>		SHILPA	<u>150MG</u>	<u>A207456 001</u>	Dec 12, 2016
<u>AB</u>			<u>500MG</u>	<u>A207456 002</u>	Dec 12, 2016
<u>AB</u>		TEVA PHARMS USA	<u>150MG</u>	<u>A091649 001</u>	Sep 16, 2013
<u>AB</u>			<u>500MG</u>	<u>A091649 002</u>	Sep 16, 2013
<u>AB</u>		TEYRO LABS	<u>150MG</u>	<u>A217237 001</u>	Oct 23, 2023
<u>AB</u>			<u>500MG</u>	<u>A217237 002</u>	Oct 23, 2023

PRESCRIPTION DRUG PRODUCT LIST

CAPECITABINE

TABLET; ORAL

XELODA

AB	+	CHEPLAPHARM	150MG	N020896	001	Apr 30, 1998
AB	+	!	500MG	N020896	002	Apr 30, 1998

CAPIVASERTIB

TABLET; ORAL

TRUQAP

	+	ASTRAZENECA	160MG	N218197	001	Nov 16, 2023
	+	!	200MG	N218197	002	Nov 16, 2023

CAPMATINIB HYDROCHLORIDE

TABLET; ORAL

TABRECTA

	+	NOVARTIS PHARM	EQ 150MG BASE	N213591	001	May 06, 2020
	+	!	EQ 200MG BASE	N213591	002	May 06, 2020

CAPSAICIN

PATCH; TOPICAL

QUTENZA

	+	AVERITAS	8%	N022395	001	Nov 16, 2009
--	---	----------	----	---------	-----	--------------

CAPTOPRIL

TABLET; ORAL

CAPTOPRIL

AB		AJANTA PHARMA LTD	12.5MG	A212809	001	Dec 13, 2019
AB			25MG	A212809	002	Dec 13, 2019
AB			50MG	A212809	003	Dec 13, 2019
AB			100MG	A212809	004	Dec 13, 2019
AB		ANDAS 5 HOLDING	12.5MG	A074677	004	May 30, 1997
AB			25MG	A074677	002	May 30, 1997
AB			50MG	A074677	001	May 30, 1997
AB			100MG	A074677	003	May 30, 1997
AB		CHANGZHOU PHARM	12.5MG	A214442	001	Jan 27, 2023
AB			25MG	A214442	002	Jan 27, 2023
AB			50MG	A214442	003	Jan 27, 2023
AB			100MG	A214442	004	Jan 27, 2023
AB		COREPHARMA	12.5MG	A074737	001	Oct 28, 1998
AB			25MG	A074737	002	Oct 28, 1998
AB			50MG	A074737	003	Oct 28, 1998
AB			100MG	A074737	004	Oct 28, 1998
AB		HIKMA INTL PHARMS	12.5MG	A074505	001	Feb 13, 1996
AB			25MG	A074505	002	Feb 13, 1996
AB			50MG	A074505	003	Feb 13, 1996
AB	!		100MG	A074505	004	Feb 13, 1996
AB		PRINSTON INC	12.5MG	A074477	001	Feb 13, 1996
AB			25MG	A074477	002	Feb 13, 1996
AB			50MG	A074477	003	Feb 13, 1996
AB			100MG	A074477	004	Feb 13, 1996
AB		WOCKHARDT	12.5MG	A074532	001	Mar 28, 1997
AB			25MG	A074532	002	Mar 28, 1997
AB			50MG	A074532	003	Mar 28, 1997
AB			100MG	A074532	004	Mar 28, 1997

CAPTOPRIL; HYDROCHLOROTHIAZIDE

TABLET; ORAL

CAPTOPRIL AND HYDROCHLOROTHIAZIDE

		RISING	25MG; 15MG	A074896	001	Dec 29, 1997
	!		25MG; 25MG	A074896	002	Dec 29, 1997
	!		50MG; 15MG	A074896	004	Dec 29, 1997
			50MG; 25MG	A074896	003	Dec 29, 1997

CARBACHOL

SOLUTION; INTRAOCULAR

MIOSTAT

	+	ALCON	0.01%	N016968	001	
--	---	-------	-------	---------	-----	--

CARBAMAZEPINE

CAPSULE, EXTENDED RELEASE; ORAL

CARBAMAZEPINE

AB		APOTEX INC	100MG	A078986	001	Nov 25, 2011
AB			200MG	A078986	002	Nov 25, 2011
AB			300MG	A078986	003	Nov 25, 2011
AB		NOSTRUM LABS INC	100MG	A076697	001	May 20, 2011
AB			200MG	A076697	002	May 20, 2011

PRESCRIPTION DRUG PRODUCT LIST

CARBAMAZEPINE

CAPSULE, EXTENDED RELEASE;ORAL

CARBAMAZEPINE

<u>AB</u>		<u>300MG</u>	<u>A076697</u>	<u>003</u>	May 20, 2011
<u>AB</u>	TARO	<u>100MG</u>	<u>A201106</u>	<u>001</u>	Jun 21, 2013
<u>AB</u>		<u>200MG</u>	<u>A201106</u>	<u>002</u>	Jun 21, 2013
<u>AB</u>		<u>300MG</u>	<u>A201106</u>	<u>003</u>	Jun 21, 2013
<u>AB</u>	TEVA PHARMS	<u>100MG</u>	<u>A078592</u>	<u>001</u>	Sep 20, 2012
<u>AB</u>		<u>200MG</u>	<u>A078592</u>	<u>002</u>	Sep 20, 2012
<u>AB</u>		<u>300MG</u>	<u>A078592</u>	<u>003</u>	Sep 20, 2012

CARBATROL

<u>AB</u>	+	TAKEDA PHARMS USA	<u>100MG</u>	<u>N020712</u>	<u>003</u>	Sep 30, 1997
<u>AB</u>	+		<u>200MG</u>	<u>N020712</u>	<u>001</u>	Sep 30, 1997
<u>AB</u>	+	!	<u>300MG</u>	<u>N020712</u>	<u>002</u>	Sep 30, 1997

EQUETRO

+	VALIDUS PHARMS	100MG	N021710	001	Dec 10, 2004
+		200MG	N021710	002	Dec 10, 2004
+	!	300MG	N021710	003	Dec 10, 2004

SUSPENSION;ORAL

CARBAMAZEPINE

<u>AB</u>	CHARTWELL RX	<u>100MG/5ML</u>	<u>A075714</u>	<u>001</u>	Jun 05, 2002
<u>AB</u>	NOVITIUM PHARMA	<u>100MG/5ML</u>	<u>A214277</u>	<u>001</u>	Oct 06, 2022

TEGRETOL

<u>AB</u>	+	NOVARTIS	<u>100MG/5ML</u>	<u>N018927</u>	<u>001</u>	Dec 18, 1987
-----------	---	----------	------------------	----------------	------------	--------------

TERIL

<u>AB</u>	TARO	<u>100MG/5ML</u>	<u>A076729</u>	<u>001</u>	Sep 20, 2004
-----------	------	------------------	----------------	------------	--------------

TABLET;ORAL

CARBAMAZEPINE

<u>AB</u>	APOTEX INC	<u>200MG</u>	<u>A075948</u>	<u>001</u>	Feb 27, 2002
<u>AB</u>	BIONPHARMA	<u>200MG</u>	<u>A218221</u>	<u>001</u>	Feb 16, 2024
<u>AB</u>	TARO	<u>200MG</u>	<u>A074649</u>	<u>001</u>	Oct 03, 1996
<u>AB</u>	TORRENT PHARMS	<u>200MG</u>	<u>A077272</u>	<u>002</u>	Dec 07, 2005
<u>AB</u>	UNICHEM	<u>200MG</u>	<u>A213284</u>	<u>001</u>	Aug 22, 2022

EPITOL

<u>AB</u>	TEVA	<u>200MG</u>	<u>A070541</u>	<u>001</u>	Sep 17, 1986
-----------	------	--------------	----------------	------------	--------------

TEGRETOL

<u>AB</u>	+	NOVARTIS	<u>200MG</u>	<u>N016608</u>	<u>001</u>	
-----------	---	----------	--------------	----------------	------------	--

CARBAMAZEPINE

BX	UMEDICA	100MG	A207798	001	Apr 15, 2020
BX		200MG	A207798	002	Apr 15, 2020

TABLET, CHEWABLE;ORAL

CARBAMAZEPINE

<u>AB</u>	TARO PHARM INDS	<u>100MG</u>	<u>A075687</u>	<u>001</u>	Oct 24, 2000
<u>AB</u>	TORRENT PHARMS	<u>100MG</u>	<u>A075712</u>	<u>001</u>	Jul 05, 2001

EPITOL

<u>AB</u>	TEVA	<u>100MG</u>	<u>A073524</u>	<u>001</u>	Jul 29, 1992
-----------	------	--------------	----------------	------------	--------------

CARBAMAZEPINE

!	TARO PHARM INDS	200MG	A075687	002	Jul 29, 2002
---	-----------------	-------	---------	-----	--------------

TABLET, EXTENDED RELEASE;ORAL

CARBAMAZEPINE

<u>AB</u>	ANBISON LAB	<u>100MG</u>	<u>A212948</u>	<u>001</u>	Sep 30, 2021
<u>AB</u>		<u>200MG</u>	<u>A212948</u>	<u>002</u>	Sep 30, 2021
<u>AB</u>		<u>400MG</u>	<u>A212948</u>	<u>003</u>	Sep 30, 2021
<u>AB</u>	CSPC OUYI	<u>100MG</u>	<u>A213311</u>	<u>001</u>	Apr 13, 2021
<u>AB</u>		<u>200MG</u>	<u>A213311</u>	<u>002</u>	Apr 13, 2021
<u>AB</u>		<u>400MG</u>	<u>A213311</u>	<u>003</u>	Apr 13, 2021
<u>AB</u>	EPIC PHARMA LLC	<u>100MG</u>	<u>A216235</u>	<u>001</u>	Mar 02, 2023
<u>AB</u>		<u>200MG</u>	<u>A216235</u>	<u>002</u>	Mar 02, 2023
<u>AB</u>		<u>400MG</u>	<u>A216235</u>	<u>003</u>	Mar 02, 2023
<u>AB</u>	NOVAST LABS	<u>100MG</u>	<u>A216404</u>	<u>001</u>	Dec 02, 2022
<u>AB</u>		<u>200MG</u>	<u>A216404</u>	<u>002</u>	Dec 02, 2022
<u>AB</u>		<u>400MG</u>	<u>A216404</u>	<u>003</u>	Dec 02, 2022
<u>AB</u>	OSMOTICA PHARM US	<u>100MG</u>	<u>A215664</u>	<u>001</u>	Oct 22, 2024
<u>AB</u>		<u>200MG</u>	<u>A215664</u>	<u>002</u>	Oct 22, 2024
<u>AB</u>		<u>400MG</u>	<u>A215664</u>	<u>003</u>	Oct 22, 2024
<u>AB</u>	TARO	<u>100MG</u>	<u>A078115</u>	<u>001</u>	Mar 31, 2009
<u>AB</u>		<u>200MG</u>	<u>A078115</u>	<u>002</u>	Mar 31, 2009
<u>AB</u>		<u>400MG</u>	<u>A078115</u>	<u>003</u>	Mar 31, 2009
<u>AB</u>	UMEDICA	<u>100MG</u>	<u>A216594</u>	<u>001</u>	Aug 18, 2022
<u>AB</u>		<u>200MG</u>	<u>A216594</u>	<u>002</u>	Aug 18, 2022
<u>AB</u>		<u>400MG</u>	<u>A216594</u>	<u>003</u>	Aug 18, 2022
<u>AB</u>	UNIQUE PHARM	<u>100MG</u>	<u>A211623</u>	<u>001</u>	Apr 24, 2020

PRESCRIPTION DRUG PRODUCT LIST

CARBAMAZEPINE

TABLET, EXTENDED RELEASE;ORAL

CARBAMAZEPINE

<u>AB</u>		<u>200MG</u>	<u>A211623</u>	<u>002</u>	Apr 24, 2020
<u>AB</u>		<u>400MG</u>	<u>A211623</u>	<u>003</u>	Apr 24, 2020
<u>AB</u>	ZHEJIANG JIUZHOU	<u>100MG</u>	<u>A215591</u>	<u>001</u>	Mar 31, 2022
<u>AB</u>		<u>200MG</u>	<u>A215591</u>	<u>002</u>	Mar 31, 2022
<u>AB</u>		<u>400MG</u>	<u>A215591</u>	<u>003</u>	Mar 31, 2022
<u>AB</u>	ZYDUS PHARMS	<u>100MG</u>	<u>A205571</u>	<u>001</u>	Feb 07, 2019
<u>AB</u>		<u>200MG</u>	<u>A205571</u>	<u>002</u>	Feb 07, 2019
<u>AB</u>		<u>400MG</u>	<u>A205571</u>	<u>003</u>	Feb 07, 2019

TEGRETOL-XR

<u>AB</u>	+	NOVARTIS	<u>100MG</u>	<u>N020234</u>	<u>001</u>	Mar 25, 1996
<u>AB</u>	+		<u>200MG</u>	<u>N020234</u>	<u>002</u>	Mar 25, 1996
<u>AB</u>	+	!	<u>400MG</u>	<u>N020234</u>	<u>003</u>	Mar 25, 1996

CARBIDOPA

TABLET;ORAL

CARBIDOPA

<u>AB</u>		ALVOGEN	<u>25MG</u>	<u>A204291</u>	<u>001</u>	Jan 08, 2016
<u>AB</u>		AUROBINDO PHARMA	<u>25MG</u>	<u>A211055</u>	<u>001</u>	Oct 21, 2019
<u>AB</u>		BEXIMCO PHARMS USA	<u>25MG</u>	<u>A217961</u>	<u>001</u>	Dec 11, 2023
<u>AB</u>		EDENBRIDGE PHARMS	<u>25MG</u>	<u>A205304</u>	<u>001</u>	Feb 17, 2016
<u>AB</u>		NOVEL LABS INC	<u>25MG</u>	<u>A204763</u>	<u>001</u>	Oct 20, 2017
<u>AB</u>		ZYDUS PHARMS	<u>25MG</u>	<u>A209910</u>	<u>001</u>	May 07, 2018

LODOSYN

<u>AB</u>	+	!	ATON	<u>25MG</u>	<u>N017830</u>	<u>001</u>
-----------	---	---	------	-------------	----------------	------------

CARBIDOPA; ENTACAPONE; LEVODOPA

TABLET;ORAL

CARBIDOPA, LEVODOPA AND ENTACAPONE

<u>AB</u>		RISING	<u>12.5MG;200MG;50MG</u>	<u>A213212</u>	<u>001</u>	Jan 25, 2022
<u>AB</u>			<u>18.75MG;200MG;75MG</u>	<u>A213212</u>	<u>002</u>	Jan 25, 2022
<u>AB</u>			<u>25MG;200MG;100MG</u>	<u>A213212</u>	<u>003</u>	Jan 25, 2022
<u>AB</u>			<u>31.25MG;200MG;125MG</u>	<u>A213212</u>	<u>004</u>	Jan 25, 2022
<u>AB</u>			<u>37.5MG;200MG;150MG</u>	<u>A213212</u>	<u>005</u>	Jan 25, 2022
<u>AB</u>			<u>50MG;200MG;200MG</u>	<u>A213212</u>	<u>006</u>	Jan 25, 2022
<u>AB</u>		SUN PHARM	<u>25MG;200MG;100MG</u>	<u>A079085</u>	<u>001</u>	May 10, 2012
<u>AB</u>			<u>37.5MG;200MG;150MG</u>	<u>A079085</u>	<u>002</u>	May 10, 2012
<u>AB</u>	+	ORION PHARMA	<u>25MG;200MG;100MG</u>	<u>N021485</u>	<u>002</u>	Jun 11, 2003
<u>AB</u>	+	ORION PHARMA	<u>31.25MG;200MG;125MG</u>	<u>N021485</u>	<u>006</u>	Aug 29, 2008
<u>AB</u>	+	ORION PHARMA	<u>37.5MG;200MG;150MG</u>	<u>N021485</u>	<u>003</u>	Jun 11, 2003
<u>AB</u>	+	ORION PHARMA	<u>50MG;200MG;200MG</u>	<u>N021485</u>	<u>004</u>	Aug 02, 2007
<u>AB</u>	+	ORION PHARMA	<u>12.5MG;200MG;50MG</u>	<u>N021485</u>	<u>001</u>	Jun 11, 2003
<u>AB</u>	+	ORION PHARMA	<u>18.75MG;200MG;75MG</u>	<u>N021485</u>	<u>005</u>	Aug 29, 2008

CARBIDOPA; LEVODOPA

CAPSULE, EXTENDED RELEASE;ORAL

CREXONT

+	IMPAX	35MG;140MG	N217186	001	Aug 07, 2024
+		52.5MG;210MG	N217186	002	Aug 07, 2024
+		70MG;280MG	N217186	003	Aug 07, 2024
+	!	87.5MG;350MG	N217186	004	Aug 07, 2024

RYTARY

+	IMPAX	23.75MG;95MG	N203312	001	Jan 07, 2015
+		36.25MG;145MG	N203312	002	Jan 07, 2015
+		48.75MG;195MG	N203312	003	Jan 07, 2015
+	!	61.25MG;245MG	N203312	004	Jan 07, 2015

SUSPENSION;ENTERAL

DUOPA

+	!	ABBVIE	4.63MG/ML;20MG/ML	N203952	001	Jan 09, 2015
---	---	--------	-------------------	---------	-----	--------------

TABLET;ORAL

CARBIDOPA AND LEVODOPA

<u>AB</u>		ACTAVIS ELIZABETH	<u>10MG;100MG</u>	<u>A074260</u>	<u>001</u>	Sep 03, 1993
<u>AB</u>			<u>25MG;100MG</u>	<u>A074260</u>	<u>002</u>	Sep 03, 1993
<u>AB</u>	!		<u>25MG;250MG</u>	<u>A074260</u>	<u>003</u>	Sep 03, 1993
<u>AB</u>		APOTEX INC	<u>10MG;100MG</u>	<u>A077120</u>	<u>001</u>	Jun 02, 2008
<u>AB</u>			<u>25MG;100MG</u>	<u>A077120</u>	<u>002</u>	Jun 02, 2008

PRESCRIPTION DRUG PRODUCT LIST

CARBIDOPA; LEVODOPA

TABLET; ORAL

CARBIDOPA AND LEVODOPA

AB		<u>25MG;250MG</u>	<u>A077120 003</u>	Jun 02, 2008
AB	AUROBINDO PHARMA LTD	<u>10MG;100MG</u>	<u>A216537 001</u>	Nov 28, 2022
AB		<u>25MG;100MG</u>	<u>A216537 002</u>	Nov 28, 2022
AB		<u>25MG;250MG</u>	<u>A216537 003</u>	Nov 28, 2022
AB	DR REDDYS LABS SA	<u>10MG;100MG</u>	<u>A073618 001</u>	Aug 28, 1992
AB		<u>25MG;100MG</u>	<u>A073589 001</u>	Aug 28, 1992
AB		<u>25MG;250MG</u>	<u>A073607 001</u>	Aug 28, 1992
AB	MYLAN	<u>10MG;100MG</u>	<u>A090324 001</u>	Sep 28, 2009
AB		<u>25MG;100MG</u>	<u>A090324 002</u>	Sep 28, 2009
AB		<u>25MG;250MG</u>	<u>A090324 003</u>	Sep 28, 2009
AB	RUBICON	<u>10MG;100MG</u>	<u>A216505 001</u>	Sep 21, 2022
AB		<u>25MG;100MG</u>	<u>A216505 002</u>	Sep 21, 2022
AB		<u>25MG;250MG</u>	<u>A216505 003</u>	Sep 21, 2022
AB	SCIEGEN PHARMS INC	<u>10MG;100MG</u>	<u>A214092 001</u>	May 07, 2021
AB		<u>25MG;100MG</u>	<u>A214092 002</u>	May 07, 2021
AB		<u>25MG;250MG</u>	<u>A214092 003</u>	May 07, 2021
AB	SUN PHARM INDS	<u>10MG;100MG</u>	<u>A078536 001</u>	Oct 28, 2008
AB		<u>25MG;100MG</u>	<u>A078536 002</u>	Oct 28, 2008
AB		<u>25MG;250MG</u>	<u>A078536 003</u>	Oct 28, 2008

SINEMET

AB	+ ORGANON	<u>10MG;100MG</u>	<u>N017555 001</u>	
AB	+	<u>25MG;100MG</u>	<u>N017555 003</u>	
AB	+	<u>25MG;250MG</u>	<u>N017555 002</u>	
	DHIVY			
	+! AVION PHARMS	25MG;100MG	N214869 001	Nov 12, 2021

TABLET, EXTENDED RELEASE; ORAL

CARBIDOPA AND LEVODOPA

AB	ACCORD HLTHCARE	<u>25MG;100MG</u>	<u>A202323 001</u>	Feb 08, 2013
AB		<u>50MG;200MG</u>	<u>A202323 002</u>	Feb 08, 2013
AB	ALEMBIC	<u>25MG;100MG</u>	<u>A210341 001</u>	Jun 05, 2019
AB		<u>50MG;200MG</u>	<u>A210341 002</u>	Jun 05, 2019
AB	APOTEX	<u>25MG;100MG</u>	<u>A076212 001</u>	Jun 16, 2004
AB		<u>50MG;200MG</u>	<u>A076212 002</u>	Jun 16, 2004
AB	IMPAX LABS	<u>25MG;100MG</u>	<u>A076521 001</u>	May 14, 2004
AB		<u>50MG;200MG</u>	<u>A076521 002</u>	May 14, 2004
AB	MYLAN	<u>25MG;100MG</u>	<u>A075091 002</u>	Apr 21, 2000
AB		<u>50MG;200MG</u>	<u>A075091 001</u>	Sep 30, 1999
AB	RUBICON	<u>25MG;100MG</u>	<u>A217482 001</u>	Jun 04, 2024
AB		<u>50MG;200MG</u>	<u>A217482 002</u>	Jun 04, 2024
AB	SCIEGEN PHARMS INC	<u>25MG;100MG</u>	<u>A214091 001</u>	Oct 05, 2021
AB	!	<u>50MG;200MG</u>	<u>A214091 002</u>	Oct 05, 2021
AB	SUN PHARM INDS	<u>25MG;100MG</u>	<u>A077828 001</u>	Aug 23, 2007
AB		<u>50MG;200MG</u>	<u>A077828 002</u>	Aug 23, 2007

TABLET, ORALLY DISINTEGRATING; ORAL

CARBIDOPA AND LEVODOPA

	SUN PHARM	10MG;100MG	A078690 001	Jul 31, 2009
		25MG;100MG	A078690 002	Jul 31, 2009
	!	25MG;250MG	A078690 003	Jul 31, 2009

CARBINOXAMINE MALEATE

SOLUTION; ORAL

CARBINOXAMINE MALEATE

!	GENUS	4MG/5ML	A040458 001	Apr 25, 2003
---	-------	---------	-------------	--------------

SUSPENSION, EXTENDED RELEASE; ORAL

KARBINAL ER

+!	AYTU	4MG/5ML	N022556 001	Mar 28, 2013
----	------	---------	-------------	--------------

TABLET; ORAL

CARBINOXAMINE MALEATE

AA	!	GENUS	<u>4MG</u>	<u>A040442 001</u>	Mar 19, 2003
AA		LEADING	<u>6MG</u>	<u>A215476 001</u>	Dec 19, 2024
AA	!	MIKART	<u>6MG</u>	<u>A207484 001</u>	May 31, 2016
AA		MISSION PHARMACAL	<u>4MG</u>	<u>A090756 001</u>	May 27, 2011

CARBOPLATIN

INJECTABLE; INTRAVENOUS

CARBOPLATIN

AP		ACCORD HLTHCARE	<u>50MG/5ML (10MG/ML)</u>	<u>A206775 001</u>	Feb 09, 2017
AP			<u>150MG/15ML (10MG/ML)</u>	<u>A206775 002</u>	Feb 09, 2017
AP			<u>450MG/45ML (10MG/ML)</u>	<u>A206775 003</u>	Feb 09, 2017
AP			<u>600MG/60ML (10MG/ML)</u>	<u>A206775 004</u>	Feb 09, 2017

PRESCRIPTION DRUG PRODUCT LIST

CARBOPLATIN

INJECTABLE; INTRAVENOUS

CARBOPLATIN

<u>AP</u>	EPIC PHARMA LLC	<u>50MG/5ML (10MG/ML)</u>	<u>A090475 001</u>	Jul 29, 2009
<u>AP</u>		<u>150MG/15ML (10MG/ML)</u>	<u>A090475 002</u>	Jul 29, 2009
<u>AP</u>		<u>450MG/45ML (10MG/ML)</u>	<u>A090475 003</u>	Jul 29, 2009
<u>AP</u>		<u>600MG/60ML (10MG/ML)</u>	<u>A091268 002</u>	Jul 28, 2010
<u>AP</u>	EUGIA PHARMA	<u>50MG/5ML (10MG/ML)</u>	<u>A205487 001</u>	Mar 28, 2016
<u>AP</u>		<u>150MG/15ML (10MG/ML)</u>	<u>A205487 002</u>	Mar 28, 2016
<u>AP</u>		<u>450MG/45ML (10MG/ML)</u>	<u>A205487 003</u>	Mar 28, 2016
<u>AP</u>		<u>600MG/60ML (10MG/ML)</u>	<u>A205487 004</u>	Aug 03, 2020
<u>AP</u>	FRESENIUS KABI USA	<u>450MG/45ML (10MG/ML)</u>	<u>A077247 003</u>	Oct 21, 2004
<u>AP</u>		<u>450MG/45ML (10MG/ML)</u>	<u>A077266 003</u>	Feb 15, 2006
<u>AP</u>		<u>600MG/60ML (10MG/ML)</u>	<u>A077266 004</u>	Feb 15, 2006
<u>AP</u>	GLAND PHARMA LTD	<u>50MG/5ML (10MG/ML)</u>	<u>A207324 001</u>	Feb 15, 2017
<u>AP</u>		<u>150MG/15ML (10MG/ML)</u>	<u>A207324 002</u>	Feb 15, 2017
<u>AP</u>		<u>450MG/45ML (10MG/ML)</u>	<u>A207324 003</u>	Feb 15, 2017
<u>AP</u>		<u>600MG/60ML (10MG/ML)</u>	<u>A207324 004</u>	Feb 15, 2017
<u>AP</u>	HIKMA	<u>50MG/5ML (10MG/ML)</u>	<u>A077244 001</u>	Oct 15, 2004
<u>AP</u>		<u>150MG/15ML (10MG/ML)</u>	<u>A077244 002</u>	Oct 15, 2004
<u>AP</u>		<u>450MG/45ML (10MG/ML)</u>	<u>A077244 003</u>	Oct 15, 2004
<u>AP</u>		<u>600MG/60ML (10MG/ML)</u>	<u>A077244 004</u>	Jan 20, 2006
<u>AP</u>	! HOSPIRA	<u>50MG/5ML (10MG/ML)</u>	<u>A076517 001</u>	Oct 14, 2004
<u>AP</u>	!	<u>150MG/15ML (10MG/ML)</u>	<u>A076517 002</u>	Oct 14, 2004
<u>AP</u>	!	<u>450MG/45ML (10MG/ML)</u>	<u>A076517 003</u>	Oct 14, 2004
<u>AP</u>	!	<u>600MG/60ML (10MG/ML)</u>	<u>A077059 001</u>	Nov 23, 2004
<u>AP</u>	NOVAST LABS	<u>50MG/5ML (10MG/ML)</u>	<u>A208487 001</u>	Apr 26, 2017
<u>AP</u>		<u>150MG/15ML (10MG/ML)</u>	<u>A208487 002</u>	Apr 26, 2017
<u>AP</u>		<u>450MG/45ML (10MG/ML)</u>	<u>A208487 003</u>	Apr 26, 2017
<u>AP</u>		<u>600MG/60ML (10MG/ML)</u>	<u>A208487 004</u>	Apr 26, 2017
<u>AP</u>	PHARMACHEMIE BV	<u>50MG/5ML (10MG/ML)</u>	<u>A077269 001</u>	Oct 14, 2004
<u>AP</u>		<u>150MG/15ML (10MG/ML)</u>	<u>A077269 002</u>	Oct 14, 2004
<u>AP</u>		<u>450MG/45ML (10MG/ML)</u>	<u>A077269 003</u>	Oct 14, 2004
<u>AP</u>		<u>600MG/60ML (10MG/ML)</u>	<u>A077269 004</u>	Dec 28, 2007
<u>AP</u>	SANDOZ	<u>50MG/5ML (10MG/ML)</u>	<u>A078280 001</u>	May 08, 2008
<u>AP</u>		<u>150MG/15ML (10MG/ML)</u>	<u>A078280 002</u>	May 08, 2008
<u>AP</u>		<u>450MG/45ML (10MG/ML)</u>	<u>A078280 003</u>	May 08, 2008
<u>AP</u>	SUN PHARM	<u>50MG/5ML (10MG/ML)</u>	<u>A077926 001</u>	Sep 19, 2008
<u>AP</u>		<u>150MG/15ML (10MG/ML)</u>	<u>A077926 002</u>	Sep 19, 2008
<u>AP</u>		<u>450MG/45ML (10MG/ML)</u>	<u>A077926 003</u>	Sep 19, 2008
<u>AP</u>	TEYRO LABS	<u>50MG/5ML (10MG/ML)</u>	<u>A077861 001</u>	Jan 18, 2007
<u>AP</u>		<u>150MG/15ML (10MG/ML)</u>	<u>A077861 002</u>	Jan 18, 2007
<u>AP</u>		<u>450MG/45ML (10MG/ML)</u>	<u>A077861 003</u>	Jan 18, 2007
<u>AP</u>		<u>600MG/60ML (10MG/ML)</u>	<u>A077861 004</u>	Jan 18, 2007
	! ACCORD HLTHCARE	<u>1GM/100ML (10MG/ML)</u>	<u>A206775 005</u>	Apr 06, 2020

CARBOPROST TROMETHAMINE

INJECTABLE; INJECTION

CARBOPROST TROMETHAMINE

<u>AP</u>	ALEMBIC	<u>EQ 0.25MG BASE/ML</u>	<u>A217198 001</u>	Jun 15, 2023
<u>AP</u>	AMNEAL	<u>EQ 0.25MG BASE/ML</u>	<u>A215337 001</u>	Jan 27, 2022
<u>AP</u>	CAPLIN	<u>EQ 0.25MG BASE/ML</u>	<u>A216882 001</u>	Feb 13, 2023
<u>AP</u>	DR REDDYS	<u>EQ 0.25MG BASE/ML</u>	<u>A211941 001</u>	Jul 02, 2019
<u>AP</u>	EUGIA PHARMA	<u>EQ 0.25MG BASE/ML</u>	<u>A216939 001</u>	May 25, 2023
<u>AP</u>	GLAND PHARMA LTD	<u>EQ 0.25MG BASE/ML</u>	<u>A217657 001</u>	Aug 07, 2023
<u>AP</u>	LONG GROVE PHARMS	<u>EQ 0.25MG BASE/ML</u>	<u>A214499 001</u>	Aug 09, 2022
<u>AP</u>	MICRO LABS	<u>EQ 0.25MG BASE/ML</u>	<u>A218013 001</u>	Apr 03, 2024
<u>AP</u>	SOLA PHARMS	<u>EQ 0.25MG BASE/ML</u>	<u>A216824 001</u>	May 19, 2023
<u>AP</u>	STERISCIENCE	<u>EQ 0.25MG BASE/ML</u>	<u>A216897 001</u>	Jul 25, 2023
	SPECLTS			
<u>AP</u>	SUNNY	<u>EQ 0.25MG BASE/ML</u>	<u>A213118 001</u>	Mar 25, 2021
	<u>HEMABATE</u>			
<u>AP</u>	+! PFIZER	<u>EQ 0.25MG BASE/ML</u>	<u>N017989 001</u>	

CARFILZOMIB

POWDER; INTRAVENOUS

CARFILZOMIB

<u>AP</u>	DR REDDYS	<u>60MG/VIAL</u>	<u>A209422 001</u>	Sep 09, 2019
	<u>KYPROLIS</u>			
<u>AP</u>	+! ONYX PHARMS AMGEN	<u>60MG/VIAL</u>	<u>N202714 001</u>	Jul 20, 2012
	+	<u>10MG/VIAL</u>	<u>N202714 003</u>	Jun 07, 2018
	+	<u>30MG/VIAL</u>	<u>N202714 002</u>	Jun 03, 2016

PRESCRIPTION DRUG PRODUCT LIST

CARGLUMIC ACID

TABLET, FOR SUSPENSION;ORAL

CARBAGLU

AB	+! RECORDATI RARE	200MG	N022562	001	Mar 18, 2010
-----------	--------------------------	--------------	----------------	------------	--------------

CARGLUMIC ACID

AB	NAVINTA LLC	200MG	A213395	001	Jun 22, 2022
-----------	-------------	--------------	----------------	------------	--------------

AB	NOVITIUM PHARMA	200MG	A213729	001	Oct 13, 2021
-----------	-----------------	--------------	----------------	------------	--------------

CARIPRAZINE HYDROCHLORIDE

CAPSULE;ORAL

VRAYLAR

+!	ABEVIE	EQ 1.5MG BASE	N204370	001	Sep 17, 2015
-----------	--------	---------------	---------	-----	--------------

+		EQ 3MG BASE	N204370	002	Sep 17, 2015
----------	--	-------------	---------	-----	--------------

+		EQ 4.5MG BASE	N204370	003	Sep 17, 2015
----------	--	---------------	---------	-----	--------------

+		EQ 6MG BASE	N204370	004	Sep 17, 2015
----------	--	-------------	---------	-----	--------------

CARISOPRODOL

TABLET;ORAL

CARISOPRODOL

AA	ACCELRX LABS	350MG	A040576	001	Jun 07, 2005
-----------	--------------	--------------	----------------	------------	--------------

AA	ALLIED	350MG	A211789	001	Oct 20, 2021
-----------	--------	--------------	----------------	------------	--------------

AA	AUROBINDO PHARMA	350MG	A040792	001	Aug 06, 2009
-----------	------------------	--------------	----------------	------------	--------------

AA	CHARTWELL RX	350MG	A040245	001	Sep 08, 1997
-----------	--------------	--------------	----------------	------------	--------------

AA		350MG	A205126	002	Jul 08, 2015
-----------	--	--------------	----------------	------------	--------------

AA	NATCO	350MG	A090988	001	Oct 28, 2014
-----------	-------	--------------	----------------	------------	--------------

AA	NOSTRUM LABS INC	350MG	A207237	002	Sep 21, 2020
-----------	------------------	--------------	----------------	------------	--------------

AA	NOVAST LABS	350MG	A040823	001	Oct 22, 2008
-----------	-------------	--------------	----------------	------------	--------------

AA	ORIENT PHARMA CO LTD	350MG	A205085	001	Oct 28, 2014
-----------	----------------------	--------------	----------------	------------	--------------

AA	OXFORD PHARMS	350MG	A040188	001	Mar 07, 1997
-----------	---------------	--------------	----------------	------------	--------------

AA	SCIEGEN PHARMS INC	350MG	A203374	001	Jan 27, 2014
-----------	--------------------	--------------	----------------	------------	--------------

AA	WANBANG BIOPHARMS	350MG	A081025	001	Apr 13, 1989
-----------	-------------------	--------------	----------------	------------	--------------

AA	WATSON LABS	350MG	A087499	001	Apr 20, 1982
-----------	-------------	--------------	----------------	------------	--------------

SOMA

AA	+ MYLAN SPECIALITY LP	350MG	N011792	001	
-----------	------------------------------	--------------	----------------	------------	--

CARISOPRODOL

AB	AUROBINDO PHARMA	250MG	A040792	002	Nov 08, 2016
-----------	------------------	--------------	----------------	------------	--------------

AB	CHARTWELL RX	250MG	A205126	001	Jul 08, 2015
-----------	--------------	--------------	----------------	------------	--------------

AB	NOSTRUM LABS INC	250MG	A207237	001	May 11, 2017
-----------	------------------	--------------	----------------	------------	--------------

SOMA

AB	+! MYLAN SPECIALITY LP	250MG	N011792	004	Sep 13, 2007
-----------	-------------------------------	--------------	----------------	------------	--------------

CARMUSTINE

IMPLANT; INTRACRANIAL

GLIADEL

+!	AZURITY	7.7MG	N020637	001	Sep 23, 1996
-----------	---------	-------	---------	-----	--------------

INJECTABLE; INJECTION

BICNU

AP	+! AVET LIFESCIENCES	100MG/VIAL	N017422	001	
-----------	-----------------------------	-------------------	----------------	------------	--

CARMUSTINE

AP	ALEMBIC	100MG/VIAL	A215730	001	Oct 20, 2023
-----------	---------	-------------------	----------------	------------	--------------

AP	AMNEAL	100MG/VIAL	A211229	001	Oct 16, 2018
-----------	--------	-------------------	----------------	------------	--------------

AP	DR REDDYS	100MG/VIAL	A213207	001	Oct 22, 2020
-----------	-----------	-------------------	----------------	------------	--------------

AP	HENGRUI PHARMA	100MG/VIAL	A211202	001	Mar 12, 2021
-----------	----------------	-------------------	----------------	------------	--------------

AP	MEITHEAL	100MG/VIAL	A213460	001	Aug 02, 2021
-----------	----------	-------------------	----------------	------------	--------------

AP	MSN	100MG/VIAL	A214814	001	May 11, 2023
-----------	-----	-------------------	----------------	------------	--------------

AP	NAVINTA LLC	100MG/VIAL	A210179	001	Sep 11, 2018
-----------	-------------	-------------------	----------------	------------	--------------

AP	PENN LIFE	100MG/VIAL	A209278	001	Apr 02, 2019
-----------	-----------	-------------------	----------------	------------	--------------

CARTEOLOL HYDROCHLORIDE

SOLUTION/DROPS;OPHTHALMIC

CARTEOLOL HYDROCHLORIDE

!	SANDOZ	1%	A075476	001	Jan 03, 2000
----------	--------	----	---------	-----	--------------

CARVEDILOL

TABLET;ORAL

CARVEDILOL

AB	AUROBINDO PHARMA	3.125MG	A078332	001	Sep 05, 2007
-----------	------------------	----------------	----------------	------------	--------------

AB		6.25MG	A078332	002	Sep 05, 2007
-----------	--	---------------	----------------	------------	--------------

AB		12.5MG	A078332	003	Sep 05, 2007
-----------	--	---------------	----------------	------------	--------------

AB		25MG	A078332	004	Sep 05, 2007
-----------	--	-------------	----------------	------------	--------------

AB	BEXIMCO USA	3.125MG	A078384	001	Sep 05, 2007
-----------	-------------	----------------	----------------	------------	--------------

AB		6.25MG	A078384	002	Sep 05, 2007
-----------	--	---------------	----------------	------------	--------------

AB		12.5MG	A078384	003	Sep 05, 2007
-----------	--	---------------	----------------	------------	--------------

AB		25MG	A078384	004	Sep 05, 2007
-----------	--	-------------	----------------	------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

CARVEDILOL

TABLET; ORAL

CARVEDILOL

<u>AB</u>	CHARTWELL MOLECULAR	<u>3.125MG</u>	<u>A077474 001</u>	Sep 05, 2007
<u>AB</u>		<u>6.25MG</u>	<u>A077474 002</u>	Sep 05, 2007
<u>AB</u>		<u>12.5MG</u>	<u>A077474 003</u>	Sep 05, 2007
<u>AB</u>		<u>25MG</u>	<u>A077474 004</u>	Sep 05, 2007
<u>AB</u>	DR REDDYS LABS LTD	<u>3.125MG</u>	<u>A076649 001</u>	Sep 05, 2007
<u>AB</u>		<u>6.25MG</u>	<u>A076649 002</u>	Sep 05, 2007
<u>AB</u>		<u>12.5MG</u>	<u>A076649 003</u>	Sep 05, 2007
<u>AB</u>		<u>25MG</u>	<u>A076649 004</u>	Sep 05, 2007
<u>AB</u>	GLENMARK PHARMS LTD	<u>3.125MG</u>	<u>A078251 001</u>	Sep 05, 2007
<u>AB</u>		<u>6.25MG</u>	<u>A078251 002</u>	Sep 05, 2007
<u>AB</u>		<u>12.5MG</u>	<u>A078251 003</u>	Sep 05, 2007
<u>AB</u>		<u>25MG</u>	<u>A078251 004</u>	Sep 05, 2007
<u>AB</u>	LUPIN	<u>3.125MG</u>	<u>A078217 001</u>	Sep 05, 2007
<u>AB</u>		<u>6.25MG</u>	<u>A078217 002</u>	Sep 05, 2007
<u>AB</u>		<u>12.5MG</u>	<u>A078217 003</u>	Sep 05, 2007
<u>AB</u>		<u>25MG</u>	<u>A078217 004</u>	Sep 05, 2007
<u>AB</u>	MLV	<u>3.125MG</u>	<u>A078786 001</u>	Dec 22, 2009
<u>AB</u>		<u>6.25MG</u>	<u>A078786 002</u>	Dec 22, 2009
<u>AB</u>		<u>12.5MG</u>	<u>A078786 003</u>	Dec 22, 2009
<u>AB</u>		<u>25MG</u>	<u>A078786 004</u>	Dec 22, 2009
<u>AB</u>	MYLAN	<u>6.25MG</u>	<u>A077316 002</u>	Sep 05, 2007
<u>AB</u>		<u>12.5MG</u>	<u>A077316 003</u>	Sep 05, 2007
<u>AB</u>		<u>25MG</u>	<u>A077316 004</u>	Sep 05, 2007
<u>AB</u>	RUBICON	<u>3.125MG</u>	<u>A078165 001</u>	Sep 05, 2007
<u>AB</u>		<u>6.25MG</u>	<u>A078165 002</u>	Sep 05, 2007
<u>AB</u>		<u>12.5MG</u>	<u>A078165 003</u>	Sep 05, 2007
<u>AB</u>		<u>25MG</u>	<u>A078165 004</u>	Sep 05, 2007
<u>AB</u>	SANDOZ	<u>3.125MG</u>	<u>A078227 001</u>	Sep 05, 2007
<u>AB</u>		<u>6.25MG</u>	<u>A078227 002</u>	Sep 05, 2007
<u>AB</u>		<u>12.5MG</u>	<u>A078227 003</u>	Sep 05, 2007
<u>AB</u>		<u>25MG</u>	<u>A078227 004</u>	Sep 05, 2007
<u>AB</u>	TARO	<u>3.125MG</u>	<u>A077780 001</u>	Sep 05, 2007
<u>AB</u>		<u>6.25MG</u>	<u>A077780 002</u>	Sep 05, 2007
<u>AB</u>		<u>12.5MG</u>	<u>A077780 003</u>	Sep 05, 2007
<u>AB</u>		<u>25MG</u>	<u>A077780 004</u>	Sep 05, 2007
<u>AB</u>	TEVA	<u>3.125MG</u>	<u>A076373 001</u>	Sep 05, 2007
<u>AB</u>		<u>6.25MG</u>	<u>A076373 002</u>	Sep 05, 2007
<u>AB</u>		<u>12.5MG</u>	<u>A076373 003</u>	Sep 05, 2007
<u>AB</u>		<u>25MG</u>	<u>A076373 004</u>	Sep 05, 2007
<u>AB</u>	ZYDUS PHARMS USA INC	<u>3.125MG</u>	<u>A077614 004</u>	Sep 05, 2007
<u>AB</u>		<u>6.25MG</u>	<u>A077614 001</u>	Sep 05, 2007
<u>AB</u>		<u>12.5MG</u>	<u>A077614 002</u>	Sep 05, 2007
<u>AB</u>		<u>25MG</u>	<u>A077614 003</u>	Sep 05, 2007

COREG

<u>AB</u>	+ WAYLIS THERAP	<u>3.125MG</u>	<u>N020297 004</u>	May 29, 1997
<u>AB</u>	+	<u>6.25MG</u>	<u>N020297 003</u>	Sep 14, 1995
<u>AB</u>	+!	<u>12.5MG</u>	<u>N020297 002</u>	Sep 14, 1995
<u>AB</u>	+	<u>25MG</u>	<u>N020297 001</u>	Sep 14, 1995

CARVEDILOL PHOSPHATE

CAPSULE, EXTENDED RELEASE; ORAL

CARVEDILOL PHOSPHATE

<u>AB</u>	IMPAX LABS INC	<u>10MG</u>	<u>A204717 001</u>	May 07, 2018
<u>AB</u>		<u>20MG</u>	<u>A204717 002</u>	May 07, 2018
<u>AB</u>		<u>40MG</u>	<u>A204717 003</u>	May 07, 2018
<u>AB</u>		<u>80MG</u>	<u>A204717 004</u>	May 07, 2018
<u>AB</u>	SUN PHARM INDUSTRIES	<u>10MG</u>	<u>A090132 001</u>	Oct 25, 2017
<u>AB</u>		<u>20MG</u>	<u>A090132 002</u>	Oct 25, 2017
<u>AB</u>	!	<u>40MG</u>	<u>A090132 003</u>	Oct 25, 2017
<u>AB</u>		<u>80MG</u>	<u>A090132 004</u>	Oct 25, 2017

CASIMERSEN

SOLUTION; INTRAVENOUS

AMONDYS 45

+	SAREPTA THERAPS INC	100MG/2ML (50MG/ML)	N213026 001	Feb 25, 2021
---	---------------------	---------------------	-------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

CASPOFUNGIN ACETATE

POWDER; INTRAVENOUS

CASPOFUNGIN ACETATE

<u>AP</u>	AREVA PHARMS	<u>50MG/VIAL</u>	<u>A211263 001</u>	Oct 01, 2021
<u>AP</u>		<u>70MG/VIAL</u>	<u>A211263 002</u>	Oct 01, 2021
<u>AP</u>	FRESENIUS KABI USA	<u>50MG/VIAL</u>	<u>N206110 001</u>	Dec 30, 2016
<u>AP</u>		<u>70MG/VIAL</u>	<u>N206110 002</u>	Dec 30, 2016
<u>AP</u>	GLAND	<u>50MG/VIAL</u>	<u>A207092 001</u>	Sep 29, 2017
<u>AP</u>		<u>70MG/VIAL</u>	<u>A207092 002</u>	Sep 29, 2017
<u>AP</u>	! HENGRUI PHARMA	<u>50MG/VIAL</u>	<u>A200833 001</u>	Jun 28, 2018
<u>AP</u>	!	<u>70MG/VIAL</u>	<u>A200833 002</u>	Jun 28, 2018

CEDAZURIDINE; DECITABINE

TABLET; ORAL

INQOVI

+! OTSUKA

100MG; 35MG

N212576 001 Jul 07, 2020

CEFACLOR

CAPSULE; ORAL

CEFACLOR

YUNG SHIN PHARM

EQ 250MG BASE

A065146 001 Jan 22, 2004

!

EQ 500MG BASE

A065146 002 Jan 22, 2004

FOR SUSPENSION; ORAL

CEFACLOR

YUNG SHIN PHARM

EQ 125MG BASE/5ML

A065412 001 Feb 17, 2012

EQ 187MG BASE/5ML

A065412 002 Feb 17, 2012

EQ 250MG BASE/5ML

A065412 003 Feb 17, 2012

!

EQ 375MG BASE/5ML

A065412 004 Feb 17, 2012

TABLET, EXTENDED RELEASE; ORAL

CEFACLOR

TEVA

EQ 375MG BASE

A065058 001 Sep 04, 2002

!

EQ 500MG BASE

A065058 002 Sep 04, 2002

CEFADROXIL/CEFADROXIL HEMIHYDRATE

CAPSULE; ORAL

CEFADROXIL

<u>AB</u>	AUROBINDO PHARMA	<u>EQ 500MG BASE</u>	<u>A065352 001</u>	Jan 25, 2007
<u>AB</u>	LUPIN	<u>EQ 500MG BASE</u>	<u>A065392 001</u>	May 29, 2007
<u>AB</u>	! TEVA PHARMS	<u>EQ 500MG BASE</u>	<u>A065282 001</u>	Jan 20, 2006

FOR SUSPENSION; ORAL

CEFADROXIL

<u>AB</u>	AUROBINDO	<u>EQ 250MG BASE/5ML</u>	<u>A065349 001</u>	Apr 25, 2013
<u>AB</u>		<u>EQ 500MG BASE/5ML</u>	<u>A065349 002</u>	Apr 25, 2013
<u>AB</u>	LUPIN	<u>EQ 250MG BASE/5ML</u>	<u>A065396 001</u>	Feb 21, 2008
<u>AB</u>	!	<u>EQ 500MG BASE/5ML</u>	<u>A065396 002</u>	Feb 21, 2008

TABLET; ORAL

CEFADROXIL

! TEVA PHARMS

EQ 1GM BASE

A062774 001 Apr 08, 1987

CEFAZOLIN SODIUM

INJECTABLE; INJECTION

CEFAZOLIN SODIUM

<u>AP</u>	ACS DOBFAR	<u>EQ 500MG BASE/VIAL</u>	<u>A065303 001</u>	Oct 22, 2008
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>A065303 002</u>	Oct 22, 2008
<u>AP</u>		<u>EQ 10GM BASE/VIAL</u>	<u>A065306 001</u>	Oct 22, 2008
<u>AP</u>	! HIKMA FARMACEUTICA	<u>EQ 500MG BASE/VIAL</u>	<u>A065047 001</u>	Sep 18, 2001
<u>AP</u>	!	<u>EQ 1GM BASE/VIAL</u>	<u>A065047 002</u>	Sep 18, 2001
<u>AP</u>		<u>EQ 10GM BASE/VIAL</u>	<u>A065143 001</u>	Oct 18, 2004
<u>AP</u>	QILU	<u>EQ 1GM BASE/VIAL</u>	<u>A203661 001</u>	Dec 28, 2015
<u>AP</u>	!	<u>EQ 10GM BASE/VIAL</u>	<u>A209217 001</u>	Oct 17, 2018
<u>AP</u>	SANDOZ	<u>EQ 500MG BASE/VIAL</u>	<u>A062831 001</u>	Dec 09, 1988
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>A062831 002</u>	Dec 09, 1988
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>A065345 001</u>	May 09, 2007
<u>AP</u>		<u>EQ 10GM BASE/VIAL</u>	<u>A062831 003</u>	Sep 25, 1992

CEFAZOLIN AND DEXTROSE

+! B BRAUN

EQ 1GM BASE/VIAL

N050779 002 Jul 27, 2000

+

EQ 2GM BASE/VIAL

N050779 003 Jan 13, 2012

+!

EQ 3GM BASE/VIAL

N050779 004 Aug 26, 2024

CEFAZOLIN SODIUM

! ACS DOBFAR

EQ 20GM BASE/VIAL

A065306 002 Aug 18, 2014

QILU

EQ 2GM BASE/VIAL

A203661 002 Mar 11, 2022

EQ 3GM BASE/VIAL

A203661 003 Jan 24, 2024

!

SAMSON MEDCL

EQ 100GM BASE/VIAL

A065141 001 Nov 29, 2006

!

EQ 300GM BASE/VIAL

A065141 002 Nov 29, 2006

PRESCRIPTION DRUG PRODUCT LIST

CEFAZOLIN SODIUM

POWDER; INTRAVENOUS

CEFAZOLIN SODIUM

+	!	HIKMA	EQ 2GM BASE/VIAL	N216109 001	Oct 07, 2022
+	!		EQ 3GM BASE/VIAL	N216109 002	Oct 07, 2022
+	!	HQ SPCLT PHARMA	EQ 2GM BASE/VIAL	N211413 001	May 08, 2023

SOLUTION; INTRAVENOUS

CEFAZOLIN IN DEXTROSE

+	!	BAXTER HLTHCARE CORP	EQ 1GM BASE/50ML (EQ 20MG BASE/ML)	N207131 002	Feb 01, 2021
+	!		EQ 2GM BASE/100ML (EQ 20MG BASE/ML)	N207131 001	Aug 07, 2015
+	!		EQ 3GM BASE/150ML (EQ 20MG BASE/ML)	N207131 003	May 31, 2024

CEFDINIR

CAPSULE; ORAL

CEFDINIR

<u>AB</u>		ALKEM LABS LTD	<u>300MG</u>	<u>A210220 001</u>	Feb 19, 2021
<u>AB</u>		ANDA REPOSITORY	<u>300MG</u>	<u>A065418 001</u>	Jul 18, 2007
<u>AB</u>		AUROBINDO PHARMA	<u>300MG</u>	<u>A065434 001</u>	Jan 07, 2008
<u>AB</u>	!	CHARTWELL RX	<u>300MG</u>	<u>A065330 001</u>	Apr 06, 2007
<u>AB</u>		LUPIN	<u>300MG</u>	<u>A065264 001</u>	May 19, 2006
<u>AB</u>		TEVA PHARMS	<u>300MG</u>	<u>A065368 001</u>	May 09, 2007

FOR SUSPENSION; ORAL

CEFDINIR

<u>AB</u>		ALKEM LABS LTD	<u>125MG/5ML</u>	<u>A210534 001</u>	Feb 19, 2021
<u>AB</u>			<u>250MG/5ML</u>	<u>A210534 002</u>	Feb 19, 2021
<u>AB</u>		ANDA REPOSITORY	<u>125MG/5ML</u>	<u>A065429 001</u>	Jul 18, 2007
<u>AB</u>			<u>250MG/5ML</u>	<u>A065429 002</u>	Jul 18, 2007
<u>AB</u>		AUROBINDO PHARMA	<u>125MG/5ML</u>	<u>A065473 001</u>	Dec 14, 2007
<u>AB</u>	!		<u>250MG/5ML</u>	<u>A065473 002</u>	Dec 14, 2007
<u>AB</u>		CHARTWELL RX	<u>125MG/5ML</u>	<u>A065337 001</u>	Apr 06, 2007
<u>AB</u>			<u>250MG/5ML</u>	<u>A065337 002</u>	Apr 06, 2007
<u>AB</u>		LUPIN	<u>125MG/5ML</u>	<u>A065259 001</u>	May 31, 2006
<u>AB</u>			<u>250MG/5ML</u>	<u>A065259 002</u>	May 07, 2007
<u>AB</u>		TEVA PHARMS	<u>125MG/5ML</u>	<u>A065332 001</u>	May 04, 2007
<u>AB</u>			<u>250MG/5ML</u>	<u>A065332 002</u>	May 04, 2007

CEFEPIME HYDROCHLORIDE

INJECTABLE; INJECTION

CEFEPIME HYDROCHLORIDE

<u>AP</u>		ACS DOBFAR	<u>EQ 1GM BASE/VIAL</u>	<u>A065441 001</u>	Mar 20, 2008
<u>AP</u>			<u>EQ 2GM BASE/VIAL</u>	<u>A065441 002</u>	Mar 20, 2008
<u>AP</u>		HIKMA	<u>EQ 1GM BASE/VIAL</u>	<u>A214402 001</u>	May 22, 2024
<u>AP</u>			<u>EQ 2GM BASE/VIAL</u>	<u>A214402 002</u>	May 22, 2024
<u>AP</u>	!	QILU	<u>EQ 1GM BASE/VIAL</u>	<u>A203704 002</u>	Feb 01, 2016
<u>AP</u>	!		<u>EQ 2GM BASE/VIAL</u>	<u>A203704 003</u>	Feb 01, 2016
<u>AP</u>		SAGENT PHARMS INC	<u>EQ 1GM BASE/VIAL</u>	<u>A091048 001</u>	Jan 04, 2017
<u>AP</u>			<u>EQ 2GM BASE/VIAL</u>	<u>A091048 002</u>	Jan 04, 2017

CEFEPIME AND DEXTROSE IN DUPLEX CONTAINER

+	!	B BRAUN	EQ 1GM BASE/VIAL	N050821 001	May 06, 2010
+	!		EQ 2GM BASE/VIAL	N050821 002	May 06, 2010

CEFEPIME HYDROCHLORIDE

!		QILU	EQ 500MG BASE/VIAL	A203704 001	Feb 01, 2016
---	--	------	--------------------	-------------	--------------

CEFEPIME IN PLASTIC CONTAINER

+	!	BAXTER HLTHCARE	EQ 1GM BASE/50ML (EQ 20MG BASE/ML)	N050817 001	Aug 05, 2008
+	!		EQ 2GM BASE/100ML (EQ 20MG BASE/ML)	N050817 002	Aug 05, 2008

POWDER; INTRAVENOUS

CEFEPIME HYDROCHLORIDE IN PLASTIC CONTAINER

!		SAMSON MEDCL	EQ 100GM BASE	A209408 001	Aug 21, 2018
---	--	--------------	---------------	-------------	--------------

CEFIDEROCOL SULFATE TOSYLATE

POWDER; INTRAVENOUS

FETROJA

+	!	SHIONOGI INC	EQ 1GM BASE/VIAL	N209445 001	Nov 14, 2019
---	---	--------------	------------------	-------------	--------------

CEFIXIME

CAPSULE; ORAL

CEFIXIME

<u>AB</u>		ALKEM LABS LTD	<u>400MG</u>	<u>A210574 001</u>	Oct 09, 2018
<u>AB</u>	+	LUPIN LTD	<u>400MG</u>	<u>N203195 001</u>	Jun 01, 2012

FOR SUSPENSION; ORAL

CEFIXIME

<u>AB</u>		ALKEM LABS LTD	<u>100MG/5ML</u>	<u>A211775 001</u>	Feb 19, 2021
<u>AB</u>			<u>200MG/5ML</u>	<u>A211775 002</u>	Feb 19, 2021

PRESCRIPTION DRUG PRODUCT LIST

CEFIXIME

FOR SUSPENSION;ORAL

CEFIXIME

<u>AB</u>	AUROBINDO PHARMA LTD	<u>100MG/5ML</u>	<u>A204835 001</u>	Apr 14, 2015
<u>AB</u>		<u>200MG/5ML</u>	<u>A204835 002</u>	Apr 14, 2015
<u>AB</u>	BELCHER	<u>100MG/5ML</u>	<u>A206938 001</u>	Feb 06, 2017
<u>AB</u>		<u>200MG/5ML</u>	<u>A206938 002</u>	Feb 06, 2017
<u>AB</u>		<u>500MG/5ML</u>	<u>A206939 001</u>	Feb 06, 2017

SUPRAX

<u>AB</u>	+! LUPIN LTD	<u>500MG/5ML</u>	<u>N202091 001</u>	Feb 20, 2013
<u>AB</u>	! LUPIN PHARMS	<u>200MG/5ML</u>	<u>A065355 001</u>	Apr 10, 2007

TABLET;ORAL

CEFIXIME

	! FDC LTD	400MG	A206358 001	Dec 17, 2024
	TABLET, CHEWABLE;ORAL			
	<u>SUPRAX</u>			
	LUPIN LTD	100MG	A065380 001	Oct 25, 2010
		150MG	A065380 002	Oct 25, 2010
	!	200MG	A065380 003	Oct 25, 2010

CEFOTETAN DISODIUM

INJECTABLE;INJECTION

CEFOTAN

<u>AP</u>	+ PAI HOLDINGS PHARM	<u>EQ 1GM BASE/VIAL</u>	<u>N050588 001</u>	Dec 27, 1985
<u>AP</u>	+	<u>EQ 2GM BASE/VIAL</u>	<u>N050588 002</u>	Dec 27, 1985

CEFOTETAN

<u>AP</u>	! FRESENIUS KABI USA	<u>EQ 1GM BASE/VIAL</u>	<u>A065374 001</u>	Aug 09, 2007
<u>AP</u>	!	<u>EQ 2GM BASE/VIAL</u>	<u>A065374 002</u>	Aug 09, 2007

CEFOXITIN SODIUM

INJECTABLE;INJECTION

CEFOXITIN

<u>AP</u>	! ACS DOBFAR	<u>EQ 1GM BASE/VIAL</u>	<u>A065414 001</u>	Jun 12, 2009
<u>AP</u>	!	<u>EQ 2GM BASE/VIAL</u>	<u>A065414 002</u>	Jun 12, 2009
<u>AP</u>	!	<u>EQ 10GM BASE/VIAL</u>	<u>A065415 001</u>	May 19, 2010
<u>AP</u>	HIKMA	<u>EQ 1GM BASE/VIAL</u>	<u>A065051 001</u>	Sep 11, 2000
<u>AP</u>		<u>EQ 2GM BASE/VIAL</u>	<u>A065051 002</u>	Sep 11, 2000
<u>AP</u>		<u>EQ 10GM BASE/VIAL</u>	<u>A065050 001</u>	Sep 11, 2000
<u>AP</u>	HIKMA FARMACEUTICA	<u>EQ 1GM BASE/VIAL</u>	<u>A065238 001</u>	Mar 12, 2010
<u>AP</u>		<u>EQ 2GM BASE/VIAL</u>	<u>A065238 002</u>	Mar 12, 2010
<u>AP</u>		<u>EQ 10GM BASE/VIAL</u>	<u>A065239 001</u>	Mar 02, 2010

CEFOXITIN AND DEXTROSE IN DUPLEX CONTAINER

<u>AP</u>	+! B BRAUN	<u>EQ 1GM BASE/VIAL</u>	<u>N065214 001</u>	Mar 10, 2006
<u>AP</u>	+!	<u>EQ 2GM BASE/VIAL</u>	<u>N065214 002</u>	Mar 10, 2006

POWDER;INTRAVENOUS

CEFOXITIN IN PLASTIC CONTAINER

	SAMSON MEDCL	EQ 100GM BASE	A200938 001	Nov 16, 2015
--	--------------	---------------	-------------	--------------

CEFPODOXIME PROXETIL

FOR SUSPENSION;ORAL

CEFPODOXIME PROXETIL

	AUROBINDO PHARMA LTD	EQ 50MG BASE/5ML	A065409 001	Jun 08, 2007
	!	EQ 100MG BASE/5ML	A065409 002	Jun 08, 2007

TABLET;ORAL

CEFPODOXIME PROXETIL

<u>AB</u>	ALKEM LABS LTD	<u>EQ 100MG BASE</u>	<u>A210568 001</u>	May 18, 2022
<u>AB</u>		<u>EQ 200MG BASE</u>	<u>A210568 002</u>	May 18, 2022
<u>AB</u>	ANDA REPOSITORY	<u>EQ 100MG BASE</u>	<u>A065388 001</u>	Nov 14, 2007
<u>AB</u>		<u>EQ 200MG BASE</u>	<u>A065388 002</u>	Nov 14, 2007
<u>AB</u>	AUROBINDO PHARMA	<u>EQ 100MG BASE</u>	<u>A065370 001</u>	Jun 11, 2007
<u>AB</u>		<u>EQ 200MG BASE</u>	<u>A065370 002</u>	Jun 11, 2007
<u>AB</u>	SANDOZ	<u>EQ 100MG BASE</u>	<u>A065462 001</u>	May 28, 2008
<u>AB</u>	!	<u>EQ 200MG BASE</u>	<u>A065462 002</u>	May 28, 2008

CEFPROZIL

FOR SUSPENSION;ORAL

CEFPROZIL

<u>AB</u>	APOTEX INC	<u>125MG/5ML</u>	<u>A065351 001</u>	Feb 29, 2012
<u>AB</u>		<u>250MG/5ML</u>	<u>A065351 002</u>	Feb 29, 2012
<u>AB</u>	AUROBINDO PHARMA	<u>125MG/5ML</u>	<u>A065381 001</u>	Jan 30, 2007
<u>AB</u>		<u>250MG/5ML</u>	<u>A065381 002</u>	Jan 30, 2007
<u>AB</u>	LUPIN	<u>125MG/5ML</u>	<u>A065261 001</u>	Dec 19, 2005
<u>AB</u>	!	<u>250MG/5ML</u>	<u>A065261 002</u>	Dec 19, 2005

PRESCRIPTION DRUG PRODUCT LIST

CEFPROZIL

TABLET; ORAL

CEFPROZIL

<u>AB</u>	ALKEM LABS LTD	<u>250MG</u>	<u>A090857 001</u>	Apr 09, 2024
<u>AB</u>		<u>500MG</u>	<u>A090857 002</u>	Apr 09, 2024
<u>AB</u>	APOTEX INC	<u>250MG</u>	<u>A065327 001</u>	Mar 26, 2008
<u>AB</u>		<u>500MG</u>	<u>A065327 002</u>	Mar 26, 2008
<u>AB</u>	AUROBINDO PHARMA	<u>250MG</u>	<u>A065340 001</u>	May 24, 2007
<u>AB</u>		<u>500MG</u>	<u>A065340 002</u>	May 24, 2007
<u>AB</u>	CHARTWELL RX	<u>250MG</u>	<u>A065235 001</u>	Nov 14, 2005
<u>AB</u>		<u>500MG</u>	<u>A065235 002</u>	Nov 14, 2005
<u>AB</u>	LUPIN	<u>250MG</u>	<u>A065276 001</u>	Dec 08, 2005
<u>AB</u>	!	<u>500MG</u>	<u>A065276 002</u>	Dec 08, 2005
<u>AB</u>	TEVA	<u>250MG</u>	<u>A065208 001</u>	Dec 06, 2005
<u>AB</u>		<u>500MG</u>	<u>A065208 002</u>	Dec 06, 2005

CEFTAROLINE FOSAMIL

POWDER; INTRAVENOUS

TEFLARO

+	!	ABBVIE	400MG/VIAL	N200327 001	Oct 29, 2010
+	!		600MG/VIAL	N200327 002	Oct 29, 2010

CEFTAZIDIME

INJECTABLE; INJECTION

CEFTAZIDIME

<u>AP</u>	!	ACS DOBFAR	<u>1GM/VIAL</u>	<u>A062640 002</u>	Nov 20, 1985
<u>AP</u>	!		<u>2GM/VIAL</u>	<u>A062640 003</u>	Nov 20, 1985
<u>AP</u>	!		<u>6GM/VIAL</u>	<u>A062640 004</u>	Feb 03, 1992

TAZICEF

<u>AP</u>		HOSPIRA	<u>1GM/VIAL</u>	<u>A062662 002</u>	Mar 06, 1986
<u>AP</u>			<u>1GM/VIAL</u>	<u>A064032 001</u>	Oct 31, 1993
<u>AP</u>			<u>2GM/VIAL</u>	<u>A062662 003</u>	Mar 06, 1986
<u>AP</u>			<u>2GM/VIAL</u>	<u>A064032 002</u>	Oct 31, 1993
<u>AP</u>			<u>6GM/VIAL</u>	<u>A062662 004</u>	Mar 06, 1986

CEFTOLOZANE SULFATE; TAZOBACTAM SODIUM

POWDER; INTRAVENOUS

ZERBAXA

+	!	CUBIST PHARMS LLC	EQ 1GM BASE/VIAL;EQ 0.5GM BASE/VIAL	N206829 001	Dec 19, 2014
---	---	-------------------	-------------------------------------	-------------	--------------

CEFTRIAZONE SODIUM

INJECTABLE; INJECTION

CEFTRIAZONE

<u>AP</u>		ACS DOBFAR	<u>EQ 500MG BASE/VIAL</u>	<u>A065329 001</u>	Jul 24, 2008
<u>AP</u>			<u>EQ 1GM BASE/VIAL</u>	<u>A065329 002</u>	Jul 24, 2008
<u>AP</u>			<u>EQ 2GM BASE/VIAL</u>	<u>A065329 003</u>	Jul 24, 2008
<u>AP</u>			<u>EQ 10GM BASE/VIAL</u>	<u>A065328 001</u>	Jul 24, 2008
<u>AP</u>		QILU	<u>EQ 10GM BASE/VIAL</u>	<u>A209218 001</u>	Oct 17, 2018
<u>AP</u>	!	SANDOZ	<u>EQ 10GM BASE/VIAL</u>	<u>A065168 001</u>	May 17, 2005
<u>AP</u>	!	SANDOZ INC	<u>EQ 1GM BASE/VIAL</u>	<u>A065204 001</u>	May 03, 2005
<u>AP</u>	!		<u>EQ 2GM BASE/VIAL</u>	<u>A065204 002</u>	May 03, 2005
<u>AP</u>		WOCKHARDT	<u>EQ 1GM BASE/VIAL</u>	<u>A065180 001</u>	May 12, 2006

CEFTRIAZONE AND DEXTROSE IN DUPLEX CONTAINER

<u>AP</u>	+	!	B BRAUN	<u>EQ 1GM BASE/VIAL</u>	<u>N050796 001</u>	Apr 20, 2005
<u>AP</u>	+	!		<u>EQ 2GM BASE/VIAL</u>	<u>N050796 002</u>	Apr 20, 2005

CEFTRIAZONE SODIUM

<u>AP</u>		ANDA REPOSITORY	<u>EQ 10GM BASE/VIAL</u>	<u>A091117 001</u>	Jan 20, 2017
<u>AP</u>		HIKMA	<u>EQ 10GM BASE/VIAL</u>	<u>A090701 001</u>	Oct 04, 2017

CEFTRIAZONE

		SAMSON MEDCL	EQ 100GM BASE/VIAL	A090057 001	Apr 25, 2014
--	--	--------------	--------------------	-------------	--------------

CEFTRIAZONE IN PLASTIC CONTAINER

!		BAXTER HLTHCARE	EQ 20MG BASE/ML	A065224 001	Aug 23, 2005
!			EQ 40MG BASE/ML	A065224 002	Aug 23, 2005

INJECTABLE; INTRAMUSCULAR, INTRAVENOUS

CEFTRIAZONE

<u>AP</u>		EPIC PHARMA LLC	<u>EQ 250MG BASE/VIAL</u>	<u>A065305 001</u>	Jan 11, 2008
<u>AP</u>			<u>EQ 500MG BASE/VIAL</u>	<u>A065305 002</u>	Jan 11, 2008
<u>AP</u>			<u>EQ 1GM BASE/VIAL</u>	<u>A065305 003</u>	Jan 11, 2008
<u>AP</u>			<u>EQ 2GM BASE/VIAL</u>	<u>A065305 004</u>	Jan 11, 2008
<u>AP</u>		HIKMA FARMACEUTICA	<u>EQ 250MG BASE/VIAL</u>	<u>A065342 001</u>	Jan 10, 2008
<u>AP</u>			<u>EQ 500MG BASE/VIAL</u>	<u>A065342 002</u>	Jan 10, 2008
<u>AP</u>			<u>EQ 1GM BASE/VIAL</u>	<u>A065342 003</u>	Jan 10, 2008
<u>AP</u>			<u>EQ 2GM BASE/VIAL</u>	<u>A065342 004</u>	Jan 10, 2008
<u>AP</u>		QILU	<u>EQ 1GM BASE/VIAL</u>	<u>A203702 003</u>	Jun 29, 2016

PRESCRIPTION DRUG PRODUCT LIST

CEFTRIAXONE SODIUM

INJECTABLE; INTRAMUSCULAR, INTRAVENOUS

CEFTRIAXONE

<u>AP</u>		<u>EQ 2GM BASE/VIAL</u>	<u>A203702 004</u>	Jun 29, 2016	
<u>AP</u>		<u>EQ 250MG BASE/VIAL</u>	<u>A203702 001</u>	Jun 29, 2016	
<u>AP</u>		<u>EQ 500MG BASE/VIAL</u>	<u>A203702 002</u>	Jun 29, 2016	
<u>AP</u>	!	SANDOZ	<u>EQ 250MG BASE/VIAL</u>	<u>A065169 001</u>	May 09, 2005
<u>AP</u>	!		<u>EQ 500MG BASE/VIAL</u>	<u>A065169 002</u>	May 09, 2005
<u>AP</u>	!		<u>EQ 1GM BASE/VIAL</u>	<u>A065169 003</u>	May 09, 2005
<u>AP</u>	!		<u>EQ 2GM BASE/VIAL</u>	<u>A065169 004</u>	May 09, 2005
<u>AP</u>		WOCKHARDT	<u>EQ 250MG BASE/VIAL</u>	<u>A065391 001</u>	Apr 12, 2007
<u>AP</u>			<u>EQ 500MG BASE/VIAL</u>	<u>A065391 002</u>	Apr 12, 2007
<u>AP</u>			<u>EQ 2GM BASE/VIAL</u>	<u>A065391 003</u>	Apr 12, 2007

CEFUROXIME AXETIL

TABLET; ORAL

CEFUROXIME AXETIL

<u>AB</u>		ALKEM LABS LTD	<u>EQ 250MG BASE</u>	<u>A065496 001</u>	Jun 07, 2010
<u>AB</u>			<u>EQ 500MG BASE</u>	<u>A065496 002</u>	Jun 07, 2010
<u>AB</u>		ANDA REPOSITORY	<u>EQ 125MG BASE</u>	<u>A065359 001</u>	Feb 15, 2008
<u>AB</u>			<u>EQ 250MG BASE</u>	<u>A065359 002</u>	Feb 15, 2008
<u>AB</u>			<u>EQ 500MG BASE</u>	<u>A065359 003</u>	Feb 15, 2008
<u>AB</u>		APOTEX	<u>EQ 250MG BASE</u>	<u>A065069 001</u>	Oct 02, 2002
<u>AB</u>			<u>EQ 500MG BASE</u>	<u>A065069 002</u>	Oct 02, 2002
<u>AB</u>		AUROBINDO PHARMA	<u>EQ 125MG BASE</u>	<u>A065308 001</u>	Mar 29, 2006
<u>AB</u>			<u>EQ 250MG BASE</u>	<u>A065308 002</u>	Mar 29, 2006
<u>AB</u>			<u>EQ 500MG BASE</u>	<u>A065308 003</u>	Mar 29, 2006
<u>AB</u>		CHARTWELL RX	<u>EQ 125MG BASE</u>	<u>A065166 001</u>	Jul 29, 2005
<u>AB</u>			<u>EQ 250MG BASE</u>	<u>A065166 002</u>	Jul 29, 2005
<u>AB</u>			<u>EQ 500MG BASE</u>	<u>A065166 003</u>	Jul 29, 2005
<u>AB</u>		LUPIN	<u>EQ 250MG BASE</u>	<u>A065135 001</u>	Jul 25, 2003
<u>AB</u>	!		<u>EQ 500MG BASE</u>	<u>A065135 002</u>	Jul 25, 2003

CEFUROXIME SODIUM

INJECTABLE; INJECTION

CEFUROXIME SODIUM

<u>AP</u>		ACS DOBFAR SPA	<u>EQ 1.5GM BASE/VIAL</u>	<u>A064125 002</u>	May 30, 1997
<u>AP</u>	!	HIKMA	<u>EQ 1.5GM BASE/VIAL</u>	<u>A065048 002</u>	Jan 09, 2004

INJECTABLE; INTRAMUSCULAR, INTRAVENOUS

CEFUROXIME SODIUM

<u>AB</u>		ACS DOBFAR SPA	<u>EQ 750MG BASE/VIAL</u>	<u>A064125 001</u>	May 30, 1997
<u>AB</u>	!	HIKMA	<u>EQ 750MG BASE/VIAL</u>	<u>A065048 001</u>	Jan 09, 2004

CELECOXIB

CAPSULE; ORAL

CELEBREX

<u>AB</u>	+	UPJOHN	<u>50MG</u>	<u>N020998 004</u>	Dec 15, 2006
<u>AB</u>	+		<u>100MG</u>	<u>N020998 001</u>	Dec 31, 1998
<u>AB</u>	+		<u>200MG</u>	<u>N020998 002</u>	Dec 31, 1998
<u>AB</u>	+	!	<u>400MG</u>	<u>N020998 003</u>	Aug 29, 2002

CELECOXIB

<u>AB</u>		ALEMBIC	<u>50MG</u>	<u>A204519 001</u>	Aug 21, 2015
<u>AB</u>			<u>100MG</u>	<u>A204519 002</u>	Aug 21, 2015
<u>AB</u>			<u>200MG</u>	<u>A204519 003</u>	Aug 21, 2015
<u>AB</u>			<u>400MG</u>	<u>A204519 004</u>	Aug 21, 2015
<u>AB</u>		APOTEX	<u>50MG</u>	<u>A204197 001</u>	Jun 02, 2015
<u>AB</u>			<u>100MG</u>	<u>A204197 002</u>	Jun 02, 2015
<u>AB</u>			<u>200MG</u>	<u>A204197 003</u>	Jun 02, 2015
<u>AB</u>		AUROBINDO PHARMA	<u>50MG</u>	<u>A206827 001</u>	Feb 01, 2016
<u>AB</u>			<u>100MG</u>	<u>A206827 002</u>	Feb 01, 2016
<u>AB</u>			<u>200MG</u>	<u>A206827 003</u>	Feb 01, 2016
<u>AB</u>			<u>400MG</u>	<u>A206827 004</u>	Feb 01, 2016
<u>AB</u>		CADILA PHARMS LTD	<u>50MG</u>	<u>A208701 001</u>	Nov 14, 2019
<u>AB</u>			<u>100MG</u>	<u>A208701 002</u>	Nov 14, 2019
<u>AB</u>			<u>200MG</u>	<u>A208701 003</u>	Nov 14, 2019
<u>AB</u>			<u>400MG</u>	<u>A208701 004</u>	Nov 14, 2019
<u>AB</u>		CIPLA	<u>50MG</u>	<u>A207446 001</u>	Sep 23, 2015
<u>AB</u>			<u>100MG</u>	<u>A207446 002</u>	Sep 23, 2015
<u>AB</u>			<u>200MG</u>	<u>A207446 003</u>	Sep 23, 2015
<u>AB</u>			<u>400MG</u>	<u>A207446 004</u>	Sep 23, 2015
<u>AB</u>		CSPC OUYI	<u>50MG</u>	<u>A210071 001</u>	Jan 23, 2018
<u>AB</u>			<u>100MG</u>	<u>A210071 002</u>	Jan 23, 2018
<u>AB</u>			<u>200MG</u>	<u>A210071 003</u>	Jan 23, 2018
<u>AB</u>		LUPIN LTD	<u>50MG</u>	<u>A202240 001</u>	Oct 29, 2014

PRESCRIPTION DRUG PRODUCT LIST

CELECOXIB

CAPSULE; ORAL

CELECOXIB

<u>AB</u>		<u>100MG</u>	<u>A202240 002</u>	Jun 09, 2015
<u>AB</u>		<u>200MG</u>	<u>A202240 003</u>	Jun 09, 2015
<u>AB</u>		<u>400MG</u>	<u>A202240 004</u>	Jun 09, 2015
<u>AB</u>	MACLEODS PHARMS LTD	<u>50MG</u>	<u>A204590 001</u>	Mar 16, 2016
<u>AB</u>		<u>100MG</u>	<u>A204590 002</u>	Mar 16, 2016
<u>AB</u>		<u>200MG</u>	<u>A204590 003</u>	Mar 16, 2016
<u>AB</u>		<u>400MG</u>	<u>A204590 004</u>	Mar 16, 2016
<u>AB</u>	MICRO LABS	<u>50MG</u>	<u>A204776 001</u>	Apr 30, 2018
<u>AB</u>		<u>100MG</u>	<u>A204776 002</u>	Apr 30, 2018
<u>AB</u>		<u>200MG</u>	<u>A204776 003</u>	Apr 30, 2018
<u>AB</u>		<u>400MG</u>	<u>A204776 004</u>	Apr 30, 2018
<u>AB</u>	NANJING	<u>200MG</u>	<u>A213598 001</u>	May 13, 2020
<u>AB</u>	QINGDAO BAHEAL PHARM	<u>50MG</u>	<u>A208856 001</u>	Aug 07, 2019
<u>AB</u>		<u>100MG</u>	<u>A208856 002</u>	Aug 07, 2019
<u>AB</u>		<u>200MG</u>	<u>A208856 003</u>	Aug 07, 2019
<u>AB</u>		<u>400MG</u>	<u>A208856 004</u>	Aug 07, 2019
<u>AB</u>	SCIEGEN PHARMS INC	<u>50MG</u>	<u>A205129 001</u>	Dec 03, 2020
<u>AB</u>		<u>100MG</u>	<u>A205129 002</u>	Dec 03, 2020
<u>AB</u>		<u>200MG</u>	<u>A205129 003</u>	Dec 03, 2020
<u>AB</u>		<u>400MG</u>	<u>A205129 004</u>	Dec 03, 2020
<u>AB</u>	TEVA	<u>50MG</u>	<u>A076898 001</u>	May 30, 2014
<u>AB</u>		<u>100MG</u>	<u>A076898 002</u>	May 30, 2014
<u>AB</u>		<u>200MG</u>	<u>A076898 003</u>	May 30, 2014
<u>AB</u>		<u>400MG</u>	<u>A076898 004</u>	May 30, 2014
<u>AB</u>	TIANJIN TIANYAO	<u>50MG</u>	<u>A207872 001</u>	Feb 25, 2020
<u>AB</u>		<u>100MG</u>	<u>A207872 002</u>	Feb 25, 2020
<u>AB</u>		<u>200MG</u>	<u>A207872 003</u>	Feb 25, 2020
<u>AB</u>		<u>400MG</u>	<u>A207872 004</u>	Feb 25, 2020
<u>AB</u>	TORRENT	<u>50MG</u>	<u>A207677 001</u>	Dec 23, 2015
<u>AB</u>		<u>100MG</u>	<u>A207677 002</u>	Dec 23, 2015
<u>AB</u>		<u>200MG</u>	<u>A207677 003</u>	Dec 23, 2015
<u>AB</u>		<u>400MG</u>	<u>A207677 004</u>	Dec 23, 2015
<u>AB</u>	UMEDICA	<u>50MG</u>	<u>A210628 001</u>	Nov 27, 2019
<u>AB</u>		<u>100MG</u>	<u>A210628 002</u>	Nov 27, 2019
<u>AB</u>		<u>200MG</u>	<u>A210628 003</u>	Nov 27, 2019
<u>AB</u>		<u>400MG</u>	<u>A210628 004</u>	Nov 27, 2019
<u>AB</u>	WATSON LABS INC	<u>50MG</u>	<u>A200562 001</u>	Feb 11, 2015
<u>AB</u>		<u>100MG</u>	<u>A200562 002</u>	Feb 11, 2015
<u>AB</u>		<u>200MG</u>	<u>A200562 003</u>	Feb 11, 2015
<u>AB</u>		<u>400MG</u>	<u>A200562 004</u>	Feb 11, 2015
<u>AB</u>	YILING	<u>50MG</u>	<u>A211412 001</u>	Mar 06, 2020
<u>AB</u>		<u>100MG</u>	<u>A211412 002</u>	Mar 06, 2020
<u>AB</u>		<u>200MG</u>	<u>A211412 003</u>	Mar 06, 2020
<u>AB</u>		<u>400MG</u>	<u>A211412 004</u>	Mar 06, 2020

SOLUTION; ORAL

ELYXYB

+! SCILEX PHARMS

25MG/ML

N212157 001 May 05, 2020

CENOBAMATE

TABLET; ORAL

XCOPRI

+! SK LIFE

12.5MG

N212839 001 Mar 10, 2020

+

25MG

N212839 002 Mar 10, 2020

+

50MG

N212839 003 Mar 10, 2020

+

100MG

N212839 004 Mar 10, 2020

+

150MG

N212839 005 Mar 10, 2020

+

200MG

N212839 006 Mar 10, 2020

CEPHALEXIN

CAPSULE; ORAL

CEPHALEXIN

<u>AB</u>	ALKEM LABS LTD	<u>EQ 250MG BASE</u>	<u>A090836 001</u>	Dec 20, 2010
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>A090836 002</u>	Dec 20, 2010
<u>AB</u>	!	<u>EQ 750MG BASE</u>	<u>A090836 004</u>	Mar 29, 2013
<u>AB</u>	ANDA REPOSITORY	<u>EQ 250MG BASE</u>	<u>A065248 001</u>	Jun 28, 2005
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>A065248 002</u>	Jun 28, 2005
<u>AB</u>	AUROBINDO PHARMA	<u>EQ 250MG BASE</u>	<u>A065253 001</u>	Nov 16, 2005
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>A065253 002</u>	Nov 16, 2005
<u>AB</u>		<u>EQ 750MG BASE</u>	<u>A065253 003</u>	Dec 06, 2023
<u>AB</u>	BELCHER PHARMS	<u>EQ 250MG BASE</u>	<u>A062713 001</u>	Jul 15, 1988

PRESCRIPTION DRUG PRODUCT LIST

CEPHALEXIN

CAPSULE; ORAL

CEPHALEXIN

<u>AB</u>		<u>EQ 500MG BASE</u>	<u>A062713 002</u>	Jul 15, 1988
<u>AB</u>	CHARTWELL RX	<u>EQ 250MG BASE</u>	<u>A065152 001</u>	Feb 24, 2005
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>A065152 002</u>	Feb 24, 2005
<u>AB</u>	LUPIN	<u>EQ 250MG BASE</u>	<u>A065229 001</u>	Nov 25, 2005
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>A065229 002</u>	Nov 25, 2005
<u>AB</u>	TEVA	<u>EQ 250MG BASE</u>	<u>A062702 001</u>	Feb 13, 1987
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>A062702 002</u>	Feb 13, 1987
	ALKEM LABS LTD	EQ 333MG BASE	A090836 003	Mar 29, 2013

FOR SUSPENSION; ORAL

CEPHALEXIN

<u>AB</u>	ALKEM LABS LTD	<u>EQ 125MG BASE/5ML</u>	<u>A210221 001</u>	Mar 26, 2019
<u>AB</u>		<u>EQ 250MG BASE/5ML</u>	<u>A210221 002</u>	Mar 26, 2019
<u>AB</u>	LUPIN	<u>EQ 125MG BASE/5ML</u>	<u>A065234 001</u>	Aug 17, 2005
<u>AB</u>		<u>EQ 250MG BASE/5ML</u>	<u>A065234 002</u>	Aug 17, 2005
<u>AB</u>	TEVA	<u>EQ 125MG BASE/5ML</u>	<u>A062703 001</u>	Feb 13, 1987
<u>AB</u>	!	<u>EQ 250MG BASE/5ML</u>	<u>A062703 002</u>	Feb 13, 1987
<u>AB</u>	YUNG SHIN PHARM	<u>EQ 125MG BASE/5ML</u>	<u>A065336 001</u>	Jul 25, 2007
<u>AB</u>		<u>EQ 250MG BASE/5ML</u>	<u>A065336 002</u>	Jul 25, 2007

TABLET; ORAL

CEPHALEXIN

<u>AB</u>	ALKEM LABS LTD	<u>EQ 250MG BASE</u>	<u>A218947 001</u>	Sep 27, 2024
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>A218947 002</u>	Sep 27, 2024
<u>AB</u>	AUROBINDO PHARMA LTD	<u>EQ 250MG BASE</u>	<u>A218817 001</u>	Sep 27, 2024
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>A218817 002</u>	Sep 27, 2024
	TEVA	EQ 250MG BASE	A063023 001	Jan 12, 1989
	!	EQ 500MG BASE	A063024 001	Jan 12, 1989

CERITINIB

TABLET; ORAL

ZYKADIA

+	!	NOVARTIS	150MG	N211225 001	Mar 18, 2019
---	---	----------	-------	-------------	--------------

CETIRIZINE HYDROCHLORIDE

SOLUTION; INTRAVENOUS

QUZYTIR

+	!	JDP	10MG/ML (10MG/ML)	N211415 001	Oct 04, 2019
---	---	-----	-------------------	-------------	--------------

SOLUTION; ORAL

CETIRIZINE HYDROCHLORIDE

<u>AA</u>	AMNEAL PHARMS	<u>5MG/5ML</u>	<u>A090766 001</u>	Oct 07, 2009	
<u>AA</u>	APOZEAL PHARMS	<u>5MG/5ML</u>	<u>A078870 001</u>	Apr 27, 2009	
<u>AA</u>	BAJAJ	<u>5MG/5ML</u>	<u>A090191 001</u>	Nov 12, 2009	
<u>AA</u>	CHARTWELL MOLECULAR	<u>5MG/5ML</u>	<u>A078876 001</u>	May 11, 2012	
<u>AA</u>	!	PADAGIS US	<u>5MG/5ML</u>	<u>A078398 001</u>	Jun 17, 2008
<u>AA</u>	PHARM ASSOC	<u>5MG/5ML</u>	<u>A078412 001</u>	Jun 18, 2008	
<u>AA</u>	TARO	<u>5MG/5ML</u>	<u>A076601 001</u>	Jun 20, 2008	

SOLUTION/DROPS; OPHTHALMIC

ZERVIAE

+	!	HARROW EYE	EQ 0.24% BASE	N208694 001	May 30, 2017
---	---	------------	---------------	-------------	--------------

CETRORELIX ACETATE

POWDER; SUBCUTANEOUS

CETRORELIX ACETATE

<u>AP</u>	GLAND	<u>EQ 0.25MG BASE/VIAL</u>	<u>A218150 001</u>	Apr 25, 2024
<u>AP</u>	LIVZON GRP	<u>EQ 0.25MG BASE/VIAL</u>	<u>A214540 001</u>	Apr 24, 2024
<u>AP</u>	QILU	<u>EQ 0.25MG BASE/VIAL</u>	<u>A217776 001</u>	Apr 16, 2024
<u>AP</u>	TEVA PHARMS INC	<u>EQ 0.25MG BASE/VIAL</u>	<u>A215737 001</u>	Aug 12, 2022

CETROTIDE

<u>AP</u>	+	!	EMD SERONO INC	<u>EQ 0.25MG BASE/VIAL</u>	<u>N021197 001</u>	Aug 11, 2000
-----------	---	---	----------------	----------------------------	--------------------	--------------

CEVIMELINE HYDROCHLORIDE

CAPSULE; ORAL

CEVIMELINE HYDROCHLORIDE

<u>AB</u>	AUROBINDO PHARMA	<u>30MG</u>	<u>A215056 001</u>	Apr 18, 2023
<u>AB</u>	BIONPHARMA	<u>30MG</u>	<u>A218290 001</u>	Nov 08, 2023
<u>AB</u>	HIKMA	<u>30MG</u>	<u>A091591 001</u>	Jul 08, 2013
<u>AB</u>	MACLEODS PHARMS LTD	<u>30MG</u>	<u>A206808 001</u>	Sep 20, 2024
<u>AB</u>	NOVEL LABS INC	<u>30MG</u>	<u>A204746 001</u>	Dec 30, 2016
<u>AB</u>	RISING	<u>30MG</u>	<u>A203775 001</u>	Jun 04, 2014
<u>AB</u>	RUBICON	<u>30MG</u>	<u>A216682 001</u>	Apr 06, 2023

PRESCRIPTION DRUG PRODUCT LIST

CEVIMELINE HYDROCHLORIDE

CAPSULE; ORAL

EVOXAC

AB	+!	COSETTE	30MG	N020989	002	Jan 11, 2000
-----------	-----------	---------	-------------	----------------	------------	--------------

CHENODIOL

TABLET; ORAL

CHENODIOL

! LGM PHARMA

250MG

A091019 001 Oct 22, 2009

CHLORAMBUCIL

TABLET; ORAL

LEUKERAN

+! WAYLIS THERAP

2MG

N010669 002

CHLORDIAZEPOXIDE HYDROCHLORIDE

CAPSULE; ORAL

CHLORDIAZEPOXIDE HYDROCHLORIDE

AB		BARR	5MG	A084768	001	
-----------	--	------	------------	----------------	------------	--

AB			10MG	A083116	001	
-----------	--	--	-------------	----------------	------------	--

AB	!		25MG	A084769	001	
-----------	----------	--	-------------	----------------	------------	--

AB		CHARTWELL RX	5MG	A084678	001	
-----------	--	--------------	------------	----------------	------------	--

AB			10MG	A084041	001	
-----------	--	--	-------------	----------------	------------	--

AB			25MG	A084679	002	
-----------	--	--	-------------	----------------	------------	--

LIBRIUM

AB	+	BAUSCH	25MG	A085475	001	
-----------	----------	--------	-------------	----------------	------------	--

AB	+	VALEANT PHARM INTL	5MG	A085461	001	
-----------	----------	--------------------	------------	----------------	------------	--

AB	+		10MG	A085472	001	
-----------	----------	--	-------------	----------------	------------	--

CHLORDIAZEPOXIDE HYDROCHLORIDE; CLIDINIUM BROMIDE

CAPSULE; ORAL

CHLORDIAZEPOXIDE HYDROCHLORIDE AND CLIDINIUM BROMIDE

AB		ALEMBIC	5MG; 2.5MG	A216969	001	Sep 15, 2023
-----------	--	---------	-------------------	----------------	------------	--------------

AB		ALKEM LABS LTD	5MG; 2.5MG	A214065	001	Apr 26, 2021
-----------	--	----------------	-------------------	----------------	------------	--------------

AB		AMNEAL	5MG; 2.5MG	A215555	001	Oct 25, 2021
-----------	--	--------	-------------------	----------------	------------	--------------

AB		AUROBINDO PHARMA LTD	5MG; 2.5MG	A216419	001	Sep 14, 2023
-----------	--	----------------------	-------------------	----------------	------------	--------------

AB		CHARTWELL RX	5MG; 2.5MG	A213530	001	Oct 20, 2020
-----------	--	--------------	-------------------	----------------	------------	--------------

AB		COREPHARMA	5MG; 2.5MG	A215453	001	Jul 29, 2022
-----------	--	------------	-------------------	----------------	------------	--------------

AB		DR REDDYS	5MG; 2.5MG	A214698	001	May 10, 2021
-----------	--	-----------	-------------------	----------------	------------	--------------

AB		MICRO LABS	5MG; 2.5MG	A215835	001	Jul 19, 2022
-----------	--	------------	-------------------	----------------	------------	--------------

AB		MISEMER	5MG; 2.5MG	A210579	001	Jul 29, 2020
-----------	--	---------	-------------------	----------------	------------	--------------

AB		NUVO PHARMS INC	5MG; 2.5MG	A211421	001	Jul 07, 2020
-----------	--	-----------------	-------------------	----------------	------------	--------------

AB		WINDER LABS LLC	5MG; 2.5MG	A212344	001	Jul 08, 2024
-----------	--	-----------------	-------------------	----------------	------------	--------------

LIBRAX

AB	+!	BAUSCH	5MG; 2.5MG	N012750	001	
-----------	-----------	--------	-------------------	----------------	------------	--

CHLORHEXIDINE GLUCONATE

SOLUTION; DENTAL

CHLORHEXIDINE GLUCONATE

AT		BAJAJ	0.12%	A075561	001	Nov 14, 2000
-----------	--	-------	--------------	----------------	------------	--------------

AT		CHARTWELL RX	0.12%	A075006	001	Mar 03, 2004
-----------	--	--------------	--------------	----------------	------------	--------------

AT		LYNE	0.12%	A074291	001	Dec 28, 1995
-----------	--	------	--------------	----------------	------------	--------------

AT		PHARM ASSOC	0.12%	A074522	001	Dec 15, 1995
-----------	--	-------------	--------------	----------------	------------	--------------

AT		XTRIUM	0.12%	A077789	001	Jun 18, 2009
-----------	--	--------	--------------	----------------	------------	--------------

PERIDEX

AT	+!	3M	0.12%	N019028	001	Aug 13, 1986
-----------	-----------	----	--------------	----------------	------------	--------------

PERIOGARD

AT		COLGATE-PALMOLIVE CO	0.12%	A203212	001	Jan 28, 2016
-----------	--	----------------------	--------------	----------------	------------	--------------

CHLOROPROCAINE HYDROCHLORIDE

GEL; OPHTHALMIC

IHEEZO

+! HARROW EYE

3%

N216227 001 Sep 27, 2022

INJECTABLE; INJECTION

CHLOROPROCAINE HYDROCHLORIDE

AP		HIKMA	2%	A040273	001	Sep 09, 1998
-----------	--	-------	-----------	----------------	------------	--------------

AP			3%	A040273	002	Sep 09, 1998
-----------	--	--	-----------	----------------	------------	--------------

NESACAINE

AP	+	FRESENIUS KABI USA	2%	N009435	002	
-----------	----------	--------------------	-----------	----------------	------------	--

NESACAINE-MPF

AP	+!	FRESENIUS KABI USA	2%	N009435	006	May 02, 1996
-----------	-----------	--------------------	-----------	----------------	------------	--------------

AP	+!		3%	N009435	007	May 02, 1996
-----------	-----------	--	-----------	----------------	------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

CHLOROQUINE PHOSPHATE

TABLET; ORAL

CHLOROQUINE PHOSPHATE

<u>AA</u>	IMPAX LABS	<u>250MG</u>	<u>A080880</u>	<u>001</u>	
<u>AA</u>		<u>500MG</u>	<u>A040516</u>	<u>001</u>	Aug 29, 2003
<u>AA</u>	! NATCO PHARMA LTD	<u>250MG</u>	<u>A091621</u>	<u>001</u>	Jan 21, 2011
<u>AA</u>	!	<u>500MG</u>	<u>A090612</u>	<u>001</u>	Jan 21, 2011
<u>AA</u>	SUVEN PHARMS	<u>500MG</u>	<u>A214756</u>	<u>001</u>	Sep 03, 2021

CHLOROTHIAZIDE

SUSPENSION; ORAL

DIURIL

+! SALIX PHARMS 250MG/5ML N011870 001

CHLOROTHIAZIDE SODIUM

INJECTABLE; INJECTION

CHLOROTHIAZIDE SODIUM

<u>AP</u>	AM REGENT	<u>EQ 500MG BASE/VIAL</u>	<u>A202561</u>	<u>001</u>	Apr 22, 2013
<u>AP</u>	FRESENIUS KABI USA	<u>EQ 500MG BASE/VIAL</u>	<u>A090896</u>	<u>001</u>	Oct 16, 2009
<u>AP</u>	GLAND PHARMA LTD	<u>EQ 500MG BASE/VIAL</u>	<u>A218630</u>	<u>001</u>	Oct 03, 2024
<u>AP</u>	! SAGENT PHARMS INC	<u>EQ 500MG BASE/VIAL</u>	<u>A202462</u>	<u>001</u>	May 29, 2015

CHLORPHENIRAMINE MALEATE; CODEINE PHOSPHATE

TABLET, EXTENDED RELEASE; ORAL

TUXARIN ER

MAINPOINTE 8MG; 54.3MG N206323 001 Jun 22, 2015

CHLORPHENIRAMINE MALEATE; HYDROCODONE BITARTRATE; PSEUDOEPHEDRINE HYDROCHLORIDE

SOLUTION; ORAL

HYDROCODONE BITARTRATE, CHLORPHENIRAMINE MALEATE AND PSEUDOEPHEDRINE HYDROCHLORIDE

! PADAGIS US 4MG/5ML; 5MG/5ML; 60MG/5ML A204627 001 Apr 29, 2014

CHLORPHENIRAMINE POLISTIREX; HYDROCODONE POLISTIREX

SUSPENSION, EXTENDED RELEASE; ORAL

HYDROCODONE POLISTIREX AND CHLORPHENIRAMINE POLISTIREX

! TRIS PHARMA INC EQ 8MG MALEATE/5ML; EQ 10MG BITARTRATE/5ML A091632 001 Oct 01, 2010

CHLORPROMAZINE HYDROCHLORIDE

CONCENTRATE; ORAL

CHLORPROMAZINE HYDROCHLORIDE

! GENUS 30MG/ML A214542 001 Jun 02, 2021
! 100MG/ML A214542 002 Jun 02, 2021

INJECTABLE; INJECTION

CHLORPROMAZINE HYDROCHLORIDE

<u>AP</u>	ASPIRO	<u>25MG/ML</u>	<u>A218272</u>	<u>001</u>	Nov 18, 2024
<u>AP</u>	DEVA HLDING	<u>25MG/ML</u>	<u>A218229</u>	<u>001</u>	Sep 24, 2024
<u>AP</u>	DR REDDYS	<u>25MG/ML</u>	<u>A080365</u>	<u>001</u>	
<u>AP</u>	EUGIA PHARMA	<u>25MG/ML</u>	<u>A211816</u>	<u>001</u>	Jul 07, 2020
<u>AP</u>	GLAND PHARMA LTD	<u>25MG/ML</u>	<u>A216911</u>	<u>001</u>	Oct 04, 2024
<u>AP</u>	+! HIKMA	<u>25MG/ML</u>	<u>A083329</u>	<u>001</u>	
<u>AP</u>	THINQ PHARM-CRO PVT	<u>25MG/ML</u>	<u>A216724</u>	<u>001</u>	Dec 10, 2024
<u>AP</u>	ZYDUS PHARMS	<u>25MG/ML</u>	<u>A217275</u>	<u>001</u>	Mar 13, 2024

TABLET; ORAL

CHLORPROMAZINE HYDROCHLORIDE

<u>AB</u>	ALEMBIC	<u>10MG</u>	<u>A217350</u>	<u>001</u>	Jul 18, 2023
<u>AB</u>		<u>25MG</u>	<u>A217350</u>	<u>002</u>	Jul 18, 2023
<u>AB</u>		<u>50MG</u>	<u>A217350</u>	<u>003</u>	Jul 18, 2023
<u>AB</u>		<u>100MG</u>	<u>A217350</u>	<u>004</u>	Jul 18, 2023
<u>AB</u>		<u>200MG</u>	<u>A217350</u>	<u>005</u>	Jul 18, 2023
<u>AB</u>	AMNEAL PHARMS CO	<u>10MG</u>	<u>A209755</u>	<u>001</u>	Sep 10, 2018
<u>AB</u>		<u>25MG</u>	<u>A209755</u>	<u>002</u>	Sep 10, 2018
<u>AB</u>		<u>50MG</u>	<u>A209755</u>	<u>003</u>	Sep 10, 2018
<u>AB</u>		<u>100MG</u>	<u>A209755</u>	<u>004</u>	Sep 10, 2018
<u>AB</u>		<u>200MG</u>	<u>A209755</u>	<u>005</u>	Sep 10, 2018
<u>AB</u>	APPCO	<u>10MG</u>	<u>A213590</u>	<u>001</u>	Aug 31, 2020
<u>AB</u>		<u>25MG</u>	<u>A213590</u>	<u>002</u>	Aug 31, 2020
<u>AB</u>		<u>50MG</u>	<u>A213590</u>	<u>003</u>	Aug 31, 2020
<u>AB</u>		<u>100MG</u>	<u>A213590</u>	<u>004</u>	Aug 31, 2020
<u>AB</u>		<u>200MG</u>	<u>A213590</u>	<u>005</u>	Aug 31, 2020
<u>AB</u>	CHARTWELL RX	<u>10MG</u>	<u>A212630</u>	<u>001</u>	Nov 29, 2021
<u>AB</u>		<u>25MG</u>	<u>A212630</u>	<u>002</u>	Nov 29, 2021
<u>AB</u>		<u>50MG</u>	<u>A212630</u>	<u>003</u>	Nov 29, 2021
<u>AB</u>		<u>100MG</u>	<u>A212630</u>	<u>004</u>	Nov 29, 2021
<u>AB</u>		<u>200MG</u>	<u>A212630</u>	<u>005</u>	Nov 29, 2021
<u>AB</u>	GLENMARK PHARMS LTD	<u>10MG</u>	<u>A212144</u>	<u>001</u>	Mar 23, 2021

PRESCRIPTION DRUG PRODUCT LIST

CHLORPROMAZINE HYDROCHLORIDE

TABLET; ORAL

CHLORPROMAZINE HYDROCHLORIDE

<u>AB</u>		<u>25MG</u>	<u>A212144</u>	<u>002</u>	Mar 23, 2021
<u>AB</u>		<u>50MG</u>	<u>A212144</u>	<u>003</u>	Mar 23, 2021
<u>AB</u>		<u>100MG</u>	<u>A212144</u>	<u>004</u>	Mar 23, 2021
<u>AB</u>		<u>200MG</u>	<u>A212144</u>	<u>005</u>	Mar 23, 2021
<u>AB</u>	LANNETT CO INC	<u>10MG</u>	<u>A212996</u>	<u>001</u>	Jan 22, 2021
<u>AB</u>		<u>25MG</u>	<u>A212996</u>	<u>002</u>	Jan 22, 2021
<u>AB</u>		<u>50MG</u>	<u>A212996</u>	<u>003</u>	Jan 22, 2021
<u>AB</u>		<u>100MG</u>	<u>A212996</u>	<u>004</u>	Jan 22, 2021
<u>AB</u>		<u>200MG</u>	<u>A212996</u>	<u>005</u>	Jan 22, 2021
<u>AB</u>	LUPIN	<u>10MG</u>	<u>A213327</u>	<u>001</u>	Jul 13, 2023
<u>AB</u>		<u>25MG</u>	<u>A213327</u>	<u>002</u>	Jul 13, 2023
<u>AB</u>		<u>50MG</u>	<u>A213327</u>	<u>003</u>	Jul 13, 2023
<u>AB</u>		<u>100MG</u>	<u>A213327</u>	<u>004</u>	Jul 13, 2023
<u>AB</u>		<u>200MG</u>	<u>A213327</u>	<u>005</u>	Jul 13, 2023
<u>AB</u>	MSN	<u>25MG</u>	<u>A214827</u>	<u>001</u>	Jan 27, 2022
<u>AB</u>		<u>50MG</u>	<u>A214827</u>	<u>002</u>	Jan 27, 2022
<u>AB</u>		<u>100MG</u>	<u>A214827</u>	<u>003</u>	Jan 27, 2022
<u>AB</u>		<u>200MG</u>	<u>A214827</u>	<u>004</u>	Jan 27, 2022
<u>AB</u>	SUN PHARM	<u>10MG</u>	<u>A214256</u>	<u>001</u>	Oct 26, 2020
<u>AB</u>		<u>25MG</u>	<u>A214256</u>	<u>002</u>	Oct 26, 2020
<u>AB</u>		<u>50MG</u>	<u>A214256</u>	<u>003</u>	Oct 26, 2020
<u>AB</u>		<u>100MG</u>	<u>A214256</u>	<u>004</u>	Oct 26, 2020
<u>AB</u>		<u>200MG</u>	<u>A214256</u>	<u>005</u>	Oct 26, 2020
<u>AB</u>	TEVA PHARMS	<u>10MG</u>	<u>A215659</u>	<u>001</u>	Oct 25, 2021
<u>AB</u>		<u>25MG</u>	<u>A215659</u>	<u>002</u>	Oct 25, 2021
<u>AB</u>		<u>50MG</u>	<u>A215659</u>	<u>003</u>	Oct 25, 2021
<u>AB</u>		<u>100MG</u>	<u>A215659</u>	<u>004</u>	Oct 25, 2021
<u>AB</u>		<u>200MG</u>	<u>A215659</u>	<u>005</u>	Oct 25, 2021
<u>AB</u>	+ UPSHER SMITH LABS	<u>10MG</u>	<u>A083386</u>	<u>001</u>	
<u>AB</u>	+!	<u>25MG</u>	<u>A084112</u>	<u>001</u>	
<u>AB</u>	+	<u>50MG</u>	<u>A084113</u>	<u>001</u>	
<u>AB</u>	+!	<u>100MG</u>	<u>A084114</u>	<u>001</u>	
<u>AB</u>	+	<u>200MG</u>	<u>A084115</u>	<u>001</u>	
<u>AB</u>	ZYDUS LIFESCIENCES	<u>10MG</u>	<u>A213368</u>	<u>001</u>	Jan 17, 2020
<u>AB</u>		<u>25MG</u>	<u>A213368</u>	<u>002</u>	Jan 17, 2020
<u>AB</u>		<u>50MG</u>	<u>A213368</u>	<u>003</u>	Jan 17, 2020
<u>AB</u>		<u>100MG</u>	<u>A213368</u>	<u>004</u>	Jan 17, 2020
<u>AB</u>		<u>200MG</u>	<u>A213368</u>	<u>005</u>	Jan 17, 2020

CHLORTHALIDONE

TABLET; ORAL

CHLORTHALIDONE

<u>AB</u>	AJANTA PHARMA LTD	<u>25MG</u>	<u>A214129</u>	<u>001</u>	Nov 27, 2020
<u>AB</u>		<u>50MG</u>	<u>A214129</u>	<u>002</u>	Nov 27, 2020
<u>AB</u>	ALEMBIC	<u>25MG</u>	<u>A216262</u>	<u>001</u>	Aug 26, 2022
<u>AB</u>		<u>50MG</u>	<u>A216262</u>	<u>002</u>	Aug 26, 2022
<u>AB</u>	ALKEM LABS LTD	<u>25MG</u>	<u>A213412</u>	<u>001</u>	Feb 11, 2020
<u>AB</u>		<u>50MG</u>	<u>A213412</u>	<u>002</u>	Feb 11, 2020
<u>AB</u>	AMNEAL PHARMS CO	<u>25MG</u>	<u>A207204</u>	<u>001</u>	Jul 01, 2019
<u>AB</u>		<u>50MG</u>	<u>A207204</u>	<u>002</u>	Jul 01, 2019
<u>AB</u>	APPCO	<u>25MG</u>	<u>A210742</u>	<u>001</u>	Oct 12, 2018
<u>AB</u>		<u>50MG</u>	<u>A210742</u>	<u>002</u>	Oct 12, 2018
<u>AB</u>	CHARTWELL RX	<u>25MG</u>	<u>A211063</u>	<u>001</u>	Feb 26, 2019
<u>AB</u>		<u>50MG</u>	<u>A211063</u>	<u>002</u>	Feb 26, 2019
<u>AB</u>	INVENTIA	<u>25MG</u>	<u>A211320</u>	<u>001</u>	Feb 09, 2022
<u>AB</u>		<u>50MG</u>	<u>A211320</u>	<u>002</u>	Feb 09, 2022
<u>AB</u>	IPCA LABS LTD	<u>25MG</u>	<u>A209162</u>	<u>001</u>	Nov 12, 2024
<u>AB</u>		<u>50MG</u>	<u>A209162</u>	<u>002</u>	Nov 12, 2024
<u>AB</u>	MANKIND PHARMA	<u>25MG</u>	<u>A215587</u>	<u>001</u>	Aug 04, 2022
<u>AB</u>		<u>50MG</u>	<u>A215587</u>	<u>002</u>	Aug 04, 2022
<u>AB</u>	+ MYLAN	<u>25MG</u>	<u>A086831</u>	<u>002</u>	
<u>AB</u>	+!	<u>50MG</u>	<u>A086831</u>	<u>001</u>	
<u>AB</u>	NOVAST LABS	<u>25MG</u>	<u>A206904</u>	<u>001</u>	Mar 30, 2017
<u>AB</u>		<u>50MG</u>	<u>A206904</u>	<u>002</u>	Mar 30, 2017
<u>AB</u>	PHARMOBEDIENT	<u>25MG</u>	<u>A212878</u>	<u>001</u>	Feb 24, 2022
<u>AB</u>		<u>50MG</u>	<u>A212875</u>	<u>001</u>	Feb 24, 2022
<u>AB</u>	SUN PHARM INDUSTRIES	<u>25MG</u>	<u>A089286</u>	<u>002</u>	Jul 21, 1986
<u>AB</u>		<u>50MG</u>	<u>A089286</u>	<u>001</u>	Jul 21, 1986
<u>AB</u>	UMEDICA	<u>25MG</u>	<u>A207222</u>	<u>001</u>	May 24, 2018

PRESCRIPTION DRUG PRODUCT LIST

CHLOROTHALIDONE

TABLET; ORAL

CHLOROTHALIDONE

<u>AB</u>		<u>50MG</u>	<u>A207222</u>	<u>002</u>	May 24, 2018
<u>AB</u>	UNICHEM	<u>25MG</u>	<u>A211627</u>	<u>001</u>	Aug 06, 2019
<u>AB</u>		<u>50MG</u>	<u>A211627</u>	<u>002</u>	Aug 06, 2019
<u>AB</u>	ZYDUS PHARMS	<u>25MG</u>	<u>A207813</u>	<u>001</u>	May 10, 2019
<u>AB</u>		<u>50MG</u>	<u>A207813</u>	<u>002</u>	May 10, 2019
THALITONE					
BX	+!	CASPER PHARMA LLC	25MG	N019574	002 Feb 12, 1992
	+		15MG	N019574	001 Dec 20, 1988

CHLORZOXAZONE

TABLET; ORAL

CHLORZOXAZONE

<u>AA</u>	AUROBINDO PHARMA LTD	<u>500MG</u>	<u>A089853</u>	<u>001</u>	May 04, 1988	
<u>AA</u>	BELCHER	<u>250MG</u>	<u>A215540</u>	<u>001</u>	Jan 24, 2023	
<u>AA</u>	COREPHARMA	<u>250MG</u>	<u>A214702</u>	<u>001</u>	Apr 10, 2023	
<u>AA</u>		<u>500MG</u>	<u>A214941</u>	<u>002</u>	Oct 20, 2022	
<u>AA</u>	DR REDDYS LABS SA	<u>500MG</u>	<u>A211849</u>	<u>002</u>	Jul 21, 2020	
<u>AA</u>	!	MIKART	<u>250MG</u>	<u>A207483</u>	<u>001</u>	Jun 24, 2016
<u>AA</u>	NOVITIUM PHARMA	<u>500MG</u>	<u>A212254</u>	<u>001</u>	Sep 12, 2019	
<u>AA</u>	RISING	<u>250MG</u>	<u>A216925</u>	<u>001</u>	Aug 09, 2023	
<u>AA</u>	SENORES PHARMS	<u>250MG</u>	<u>A215158</u>	<u>001</u>	Jul 29, 2021	
<u>AA</u>		<u>500MG</u>	<u>A210961</u>	<u>001</u>	Jan 22, 2024	
<u>AA</u>	!	WATSON LABS	<u>500MG</u>	<u>A089859</u>	<u>001</u>	May 04, 1988
<u>AB</u>	AUROBINDO PHARMA LTD	<u>375MG</u>	<u>A089853</u>	<u>002</u>	Aug 17, 2023	
<u>AB</u>		<u>750MG</u>	<u>A089853</u>	<u>003</u>	Aug 17, 2023	
<u>AB</u>	COREPHARMA	<u>375MG</u>	<u>A214941</u>	<u>001</u>	Oct 20, 2022	
<u>AB</u>		<u>750MG</u>	<u>A214941</u>	<u>003</u>	Oct 20, 2022	
<u>AB</u>	DR REDDYS LABS SA	<u>375MG</u>	<u>A211849</u>	<u>001</u>	Jul 21, 2020	
<u>AB</u>		<u>750MG</u>	<u>A211849</u>	<u>003</u>	Jul 21, 2020	
<u>AB</u>	ENDO OPERATIONS	<u>375MG</u>	<u>A212743</u>	<u>002</u>	Apr 29, 2021	
<u>AB</u>		<u>750MG</u>	<u>A212743</u>	<u>001</u>	Nov 02, 2020	
<u>AB</u>	!	MIKART	<u>375MG</u>	<u>A040861</u>	<u>001</u>	Jun 01, 2010
<u>AB</u>	!		<u>750MG</u>	<u>A040861</u>	<u>002</u>	Jun 01, 2010
<u>AB</u>	NOVITIUM PHARMA	<u>375MG</u>	<u>A212253</u>	<u>001</u>	Nov 27, 2019	
<u>AB</u>		<u>750MG</u>	<u>A212253</u>	<u>002</u>	Nov 27, 2019	
<u>AB</u>	RISING	<u>375MG</u>	<u>A213126</u>	<u>001</u>	Apr 05, 2022	
<u>AB</u>		<u>500MG</u>	<u>A213126</u>	<u>002</u>	Apr 05, 2022	
<u>AB</u>		<u>750MG</u>	<u>A213126</u>	<u>003</u>	Apr 05, 2022	
<u>AB</u>	UPSHER SMITH LABS	<u>375MG</u>	<u>A212047</u>	<u>001</u>	Jan 27, 2023	
<u>AB</u>		<u>750MG</u>	<u>A212047</u>	<u>002</u>	Jan 27, 2023	

CHOLESTYRAMINE

POWDER; ORAL

CHOLESTYRAMINE

<u>AB</u>	AJANTA PHARMA LTD	<u>EQ 4GM RESIN/PACKET</u>	<u>A211119</u>	<u>001</u>	Apr 06, 2020	
<u>AB</u>		<u>EQ 4GM RESIN/SCOOPFUL</u>	<u>A211119</u>	<u>002</u>	Apr 06, 2020	
<u>AB</u>	ALKEM LABS LTD	<u>EQ 4GM RESIN/PACKET</u>	<u>A211856</u>	<u>001</u>	Oct 19, 2021	
<u>AB</u>		<u>EQ 4GM RESIN/SCOOPFUL</u>	<u>A211856</u>	<u>002</u>	Oct 19, 2021	
<u>AB</u>	ENDO OPERATIONS	<u>EQ 4GM RESIN/PACKET</u>	<u>A077204</u>	<u>001</u>	Aug 26, 2005	
<u>AB</u>		<u>EQ 4GM RESIN/SCOOPFUL</u>	<u>A077204</u>	<u>002</u>	Aug 26, 2005	
<u>AB</u>	!	EPIC PHARMA LLC	<u>EQ 4GM RESIN/PACKET</u>	<u>A074557</u>	<u>001</u>	Aug 15, 1996
<u>AB</u>		<u>EQ 4GM RESIN/SCOOPFUL</u>	<u>A074557</u>	<u>002</u>	Aug 15, 1996	
<u>AB</u>	MACLEODS PHARMS LTD	<u>EQ 4GM RESIN/PACKET</u>	<u>A207294</u>	<u>001</u>	Nov 21, 2024	
<u>AB</u>		<u>EQ 4GM RESIN/SCOOPFUL</u>	<u>A207294</u>	<u>002</u>	Nov 21, 2024	
<u>AB</u>	TAGI	<u>EQ 4GM RESIN/PACKET</u>	<u>A209597</u>	<u>001</u>	Mar 09, 2021	
<u>AB</u>		<u>EQ 4GM RESIN/SCOOPFUL</u>	<u>A209597</u>	<u>002</u>	Mar 09, 2021	
<u>AB</u>	ZYDUS PHARMS	<u>EQ 4GM RESIN/SCOOPFUL</u>	<u>A202901</u>	<u>001</u>	Jul 02, 2018	

CHOLESTYRAMINE LIGHT

<u>AB</u>	ALKEM LABS LTD	<u>EQ 4GM RESIN/PACKET</u>	<u>A211799</u>	<u>001</u>	Oct 19, 2021	
<u>AB</u>		<u>EQ 4GM RESIN/SCOOPFUL</u>	<u>A211799</u>	<u>002</u>	Oct 19, 2021	
<u>AB</u>	ENDO OPERATIONS	<u>EQ 4GM RESIN/PACKET</u>	<u>A077203</u>	<u>001</u>	Aug 26, 2005	
<u>AB</u>		<u>EQ 4GM RESIN/SCOOPFUL</u>	<u>A077203</u>	<u>002</u>	Aug 26, 2005	
<u>AB</u>	!	EPIC PHARMA LLC	<u>EQ 4GM RESIN/PACKET</u>	<u>A074558</u>	<u>001</u>	Aug 15, 1996
<u>AB</u>		<u>EQ 4GM RESIN/SCOOPFUL</u>	<u>A074558</u>	<u>002</u>	Aug 15, 1996	
<u>AB</u>	MACLEODS PHARMS LTD	<u>EQ 4GM RESIN/SCOOPFUL</u>	<u>A209109</u>	<u>001</u>	Oct 31, 2024	
<u>AB</u>		<u>EQ 4GM RESIN/PACKET</u>	<u>A209109</u>	<u>002</u>	Oct 31, 2024	
<u>AB</u>	TAGI	<u>EQ 4GM RESIN/PACKET</u>	<u>A209599</u>	<u>001</u>	Nov 12, 2020	
<u>AB</u>		<u>EQ 4GM RESIN/SCOOPFUL</u>	<u>A209599</u>	<u>002</u>	Nov 12, 2020	
<u>AB</u>	ZYDUS PHARMS	<u>EQ 4GM RESIN/SCOOPFUL</u>	<u>A202902</u>	<u>001</u>	Apr 25, 2017	

PRESCRIPTION DRUG PRODUCT LIST

CHOLESTYRAMINE

POWDER; ORAL

LOCHOLEST

AB	CHARTWELL RX	EQ 4GM RESIN/PACKET	A074561 001	Aug 15, 1996
AB		EQ 4GM RESIN/SCOOPFUL	A074561 002	Aug 15, 1996

LOCHOLEST LIGHT

AB	CHARTWELL RX	EQ 4GM RESIN/PACKET	A074562 001	Aug 15, 1996
AB		EQ 4GM RESIN/SCOOPFUL	A074562 002	Aug 15, 1996

PREVALITE

AB	UPSHER SMITH LABS	EQ 4GM RESIN/PACKET	A073263 001	Feb 22, 1996
AB		EQ 4GM RESIN/SCOOPFUL	A073263 002	Oct 30, 1997

CHOLIC ACID

CAPSULE; ORAL

CHOLBAM

	+ MIRUM	50MG	N205750 001	Mar 17, 2015
	+!	250MG	N205750 002	Mar 17, 2015

CHOLINE C-11

INJECTABLE; INTRAVENOUS

CHOLINE C-11

AP	DECATUR	4-33.1mCi/ML	A206319 001	Nov 13, 2015
AP	+! MCPRF	4-33.1mCi/ML	N203155 001	Sep 12, 2012

CHOLINE FENOFIBRATE

CAPSULE, DELAYED RELEASE; ORAL

FENOFIBRIC ACID

AB	ACTAVIS ELIZABETH	EQ 45MG FENOFIBRIC ACID	A200920 001	Oct 07, 2015
AB		EQ 135MG FENOFIBRIC ACID	A200920 002	Oct 07, 2015
AB	ALEMbic	EQ 45MG FENOFIBRIC ACID	A208705 001	May 12, 2017
AB		EQ 135MG FENOFIBRIC ACID	A208705 002	May 12, 2017
AB	AUROBINDO PHARMA LTD	EQ 45MG FENOFIBRIC ACID	A212598 001	Jul 25, 2019
AB		EQ 135MG FENOFIBRIC ACID	A212598 002	Jul 25, 2019
AB	CHARTWELL RX	EQ 45MG FENOFIBRIC ACID	A200750 001	Dec 04, 2013
AB		EQ 135MG FENOFIBRIC ACID	A200750 002	Dec 04, 2013
AB	IMPAX LABS INC	EQ 45MG FENOFIBRIC ACID	A200264 001	Sep 07, 2016
AB		EQ 135MG FENOFIBRIC ACID	A200264 002	Sep 07, 2016
AB	MICRO LABS	EQ 45MG FENOFIBRIC ACID	A213450 001	Jun 16, 2020
AB		EQ 135MG FENOFIBRIC ACID	A213450 002	Jun 16, 2020
AB	RISING	EQ 45MG FENOFIBRIC ACID	A211626 001	Jul 18, 2019
AB		EQ 135MG FENOFIBRIC ACID	A211626 002	Jul 18, 2019
AB	YICHANG HUMANWELL	EQ 45MG FENOFIBRIC ACID	A212562 001	Dec 23, 2020
AB		EQ 135MG FENOFIBRIC ACID	A212562 002	Dec 23, 2020

TRILIPIX

AB	+ ABBVIE	EQ 45MG FENOFIBRIC ACID	N022224 001	Dec 15, 2008
AB	+!	EQ 135MG FENOFIBRIC ACID	N022224 002	Dec 15, 2008

CHROMIC CHLORIDE

INJECTABLE; INJECTION

CHROMIC CHLORIDE

AP	RK PHARMA	EQ 0.004MG CHROMIUM/ML	A218538 001	Nov 18, 2024
-----------	-----------	-------------------------------	--------------------	--------------

CHROMIC CHLORIDE IN PLASTIC CONTAINER

AP	+! HOSPIRA	EQ 0.004MG CHROMIUM/ML	N018961 001	Jun 26, 1986
-----------	------------	-------------------------------	--------------------	--------------

CICLESONIDE

AEROSOL, METERED; INHALATION

ALVESCO

	+! COVIS	0.08MG/INH	N021658 002	Jan 10, 2008
	+!	0.16MG/INH	N021658 003	Jan 10, 2008

SPRAY, METERED; NASAL

OMNARIS

	+! COVIS	0.05MG/SPRAY	N022004 001	Oct 20, 2006
--	----------	--------------	-------------	--------------

CICLOPIROX

CREAM; TOPICAL

CICLOPIROX

AB	COSETTE	0.77%	A078463 001	Dec 20, 2010
AB	FOUGERA PHARMS	0.77%	A076435 001	Dec 29, 2004
AB	GLENMARK PHARMS	0.77%	A090273 001	Nov 10, 2009
AB	PADAGIS ISRAEL	0.77%	A077364 001	Mar 03, 2006
AB	TARO	0.77%	A076790 001	Apr 12, 2005

LOPROX

AB	+! MEDIMETRIKS PHARMS	0.77%	N018748 001	Dec 30, 1982
-----------	-----------------------	--------------	--------------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

CICLOPIROX

GEL; TOPICAL

CICLOPIROX

<u>AB</u>	FOUGERA PHARMS	<u>0.77%</u>	<u>A077896</u>	<u>001</u>	Jun 10, 2008
<u>AB</u>	! GLENMARK PHARMS INC	<u>0.77%</u>	<u>A091595</u>	<u>001</u>	Feb 29, 2012
<u>AB</u>	PADAGIS US	<u>0.77%</u>	<u>A078266</u>	<u>001</u>	Jan 07, 2009

SHAMPOO; TOPICAL

CICLOPIROX

<u>AT</u>	ACTAVIS MID ATLANTIC	<u>1%</u>	<u>A090490</u>	<u>001</u>	Nov 24, 2009
<u>AT</u>	FOUGERA PHARMS	<u>1%</u>	<u>A090146</u>	<u>001</u>	May 25, 2010
<u>AT</u>	PADAGIS US	<u>1%</u>	<u>A078594</u>	<u>001</u>	Feb 16, 2010
<u>AT</u>	TARO	<u>1%</u>	<u>A090269</u>	<u>001</u>	Feb 23, 2011

LOPROX

<u>AT</u>	+! BAUSCH	<u>1%</u>	<u>N021159</u>	<u>001</u>	Feb 28, 2003
-----------	-----------	-----------	----------------	------------	--------------

SOLUTION; TOPICAL

CICLOPIROX

<u>AT</u>	ACELLA	<u>8%</u>	<u>A078172</u>	<u>001</u>	Sep 18, 2007
<u>AT</u>	CHARTWELL RX	<u>8%</u>	<u>A078046</u>	<u>001</u>	Sep 18, 2007
<u>AT</u>	ENCUBE	<u>8%</u>	<u>A077687</u>	<u>001</u>	Sep 18, 2007
<u>AT</u>	! PADAGIS US	<u>8%</u>	<u>A077623</u>	<u>001</u>	Sep 18, 2007
<u>AT</u>	RISING	<u>8%</u>	<u>A078124</u>	<u>001</u>	Sep 18, 2007
<u>AT</u>	TARO PHARM INDS	<u>8%</u>	<u>A078144</u>	<u>001</u>	Sep 18, 2007

SUSPENSION; TOPICAL

CICLOPIROX

<u>AB</u>	FOUGERA PHARMS	<u>0.77%</u>	<u>A076422</u>	<u>001</u>	Aug 06, 2004
<u>AB</u>	PADAGIS ISRAEL	<u>0.77%</u>	<u>A077676</u>	<u>001</u>	Dec 15, 2006
<u>AB</u>	TARO	<u>0.77%</u>	<u>A077092</u>	<u>001</u>	Aug 10, 2005

LOPROX

<u>AB</u>	+! MEDIMETRIKS PHARMS	<u>0.77%</u>	<u>N019824</u>	<u>001</u>	Dec 30, 1988
-----------	-----------------------	--------------	----------------	------------	--------------

CIDOFOVIR

INJECTABLE; INJECTION

CIDOFOVIR

<u>AP</u>	AVET LIFESCIENCES	<u>EQ 75MG BASE/ML</u>	<u>A202501</u>	<u>001</u>	Jul 26, 2012
<u>AP</u>	! MYLAN INSTITUTIONAL	<u>EQ 75MG BASE/ML</u>	<u>A201276</u>	<u>001</u>	Jun 27, 2012

CILASTATIN SODIUM; IMIPENEM

POWDER; INTRAVENOUS

IMIPENEM AND CILASTATIN

<u>AP</u>	ACS DOBFAR	<u>EQ 500MG BASE/VIAL; 500MG/VIAL</u>	<u>A090577</u>	<u>002</u>	Dec 21, 2011
<u>AP</u>	HQ SPCLT PHARMA	<u>EQ 500MG BASE/VIAL; 500MG/VIAL</u>	<u>A207594</u>	<u>001</u>	Dec 12, 2019

PRIMAXIN

<u>AP</u>	+! MERCK	<u>EQ 500MG BASE/VIAL; 500MG/VIAL</u>	<u>N050587</u>	<u>002</u>	Nov 26, 1985
	IMIPENEM AND CILASTATIN				
	! ACS DOBFAR	<u>EQ 250MG BASE/VIAL; 250MG/VIAL</u>	<u>A090577</u>	<u>001</u>	Dec 21, 2011

CILASTATIN SODIUM; IMIPENEM; RELEBACTAM

POWDER; INTRAVENOUS

RECARBRIO

	+! MSD MERCK CO	<u>EQ 500MG BASE/VIAL; 500MG/VIAL; 250MG/VIAL</u>	<u>N212819</u>	<u>001</u>	Jul 16, 2019
--	-----------------	---	----------------	------------	--------------

CILOSTAZOL

TABLET; ORAL

CILOSTAZOL

<u>AB</u>	APOTEX INC	<u>50MG</u>	<u>A077030</u>	<u>001</u>	Dec 10, 2004
<u>AB</u>		<u>100MG</u>	<u>A077030</u>	<u>002</u>	Dec 10, 2004
<u>AB</u>	CHARTWELL RX	<u>50MG</u>	<u>A077831</u>	<u>002</u>	Sep 24, 2012
<u>AB</u>		<u>100MG</u>	<u>A077831</u>	<u>001</u>	Sep 24, 2012
<u>AB</u>	EPIC PHARMA LLC	<u>50MG</u>	<u>A077022</u>	<u>002</u>	Mar 11, 2005
<u>AB</u>		<u>100MG</u>	<u>A077022</u>	<u>001</u>	Nov 23, 2004
<u>AB</u>	SLATE RUN PHARMA	<u>50MG</u>	<u>A077208</u>	<u>002</u>	Mar 29, 2006
<u>AB</u>		<u>100MG</u>	<u>A077208</u>	<u>001</u>	Mar 29, 2006
<u>AB</u>	! TEVA	<u>50MG</u>	<u>A077027</u>	<u>001</u>	Nov 24, 2004
<u>AB</u>	!	<u>100MG</u>	<u>A077027</u>	<u>002</u>	Nov 24, 2004

CIMETIDINE

TABLET; ORAL

CIMETIDINE

<u>AB</u>	AUROBINDO PHARMA LTD	<u>200MG</u>	<u>A218997</u>	<u>001</u>	Nov 12, 2024
<u>AB</u>		<u>300MG</u>	<u>A218997</u>	<u>002</u>	Nov 12, 2024
<u>AB</u>		<u>400MG</u>	<u>A218997</u>	<u>003</u>	Nov 12, 2024
<u>AB</u>		<u>800MG</u>	<u>A218997</u>	<u>004</u>	Nov 12, 2024

PRESCRIPTION DRUG PRODUCT LIST

CIMETIDINE

TABLET; ORAL

CIMETIDINE

<u>AB</u>	BIONPHARMA	<u>200MG</u>	<u>A218220 001</u>	Nov 07, 2024
<u>AB</u>		<u>300MG</u>	<u>A218220 002</u>	Nov 07, 2024
<u>AB</u>		<u>400MG</u>	<u>A218220 003</u>	Nov 07, 2024
<u>AB</u>		<u>800MG</u>	<u>A218220 004</u>	Nov 07, 2024
<u>AB</u>	CHARTWELL MOLECULES	<u>200MG</u>	<u>A074329 002</u>	May 17, 1994
<u>AB</u>		<u>300MG</u>	<u>A074329 003</u>	May 17, 1994
<u>AB</u>		<u>400MG</u>	<u>A074329 004</u>	May 17, 1994
<u>AB</u>		<u>800MG</u>	<u>A074329 001</u>	May 17, 1994
<u>AB</u>	MYLAN	<u>200MG</u>	<u>A074246 001</u>	May 17, 1994
<u>AB</u>		<u>300MG</u>	<u>A074246 002</u>	May 17, 1994
<u>AB</u>		<u>400MG</u>	<u>A074246 003</u>	May 17, 1994
<u>AB</u>	!	<u>800MG</u>	<u>A074246 004</u>	May 17, 1994
<u>AB</u>	TEVA	<u>200MG</u>	<u>A074151 001</u>	May 17, 1994
<u>AB</u>		<u>300MG</u>	<u>A074151 002</u>	May 17, 1994
<u>AB</u>		<u>400MG</u>	<u>A074151 003</u>	May 17, 1994
<u>AB</u>		<u>800MG</u>	<u>A074463 001</u>	May 17, 1994

CIMETIDINE HYDROCHLORIDE

SOLUTION; ORAL

CIMETIDINE HYDROCHLORIDE

<u>AA</u>	CHARTWELL MOLECULAR	<u>EQ 300MG BASE/5ML</u>	<u>A074251 001</u>	Dec 22, 1994
<u>AA</u>	!	<u>EQ 300MG BASE/5ML</u>	<u>A074664 001</u>	Oct 28, 1997

CINACALCET HYDROCHLORIDE

TABLET; ORAL

CINACALCET HYDROCHLORIDE

<u>AB</u>	ACCORD HLTHCARE	<u>EQ 30MG BASE</u>	<u>A211892 001</u>	May 15, 2020
<u>AB</u>		<u>EQ 60MG BASE</u>	<u>A211892 002</u>	May 15, 2020
<u>AB</u>		<u>EQ 90MG BASE</u>	<u>A211892 003</u>	May 15, 2020
<u>AB</u>	ALKEM LABS LTD	<u>EQ 30MG BASE</u>	<u>A210570 001</u>	May 17, 2019
<u>AB</u>		<u>EQ 60MG BASE</u>	<u>A210570 002</u>	May 17, 2019
<u>AB</u>		<u>EQ 90MG BASE</u>	<u>A210570 003</u>	May 17, 2019
<u>AB</u>	AUROBINDO PHARMA LTD	<u>EQ 30MG BASE</u>	<u>A206125 001</u>	Mar 08, 2018
<u>AB</u>		<u>EQ 60MG BASE</u>	<u>A206125 002</u>	Mar 08, 2018
<u>AB</u>		<u>EQ 90MG BASE</u>	<u>A206125 003</u>	Mar 08, 2018
<u>AB</u>	CHARTWELL RX	<u>EQ 30MG BASE</u>	<u>A213325 001</u>	May 18, 2020
<u>AB</u>		<u>EQ 60MG BASE</u>	<u>A213325 002</u>	May 18, 2020
<u>AB</u>		<u>EQ 90MG BASE</u>	<u>A213325 003</u>	May 18, 2020
<u>AB</u>	CIPLA	<u>EQ 30MG BASE</u>	<u>A208915 001</u>	Mar 08, 2018
<u>AB</u>		<u>EQ 60MG BASE</u>	<u>A208915 002</u>	Mar 08, 2018
<u>AB</u>		<u>EQ 90MG BASE</u>	<u>A208915 003</u>	Mar 08, 2018
<u>AB</u>	DR REDDYS	<u>EQ 30MG BASE</u>	<u>A208368 001</u>	Sep 18, 2020
<u>AB</u>		<u>EQ 60MG BASE</u>	<u>A208368 002</u>	Sep 18, 2020
<u>AB</u>		<u>EQ 90MG BASE</u>	<u>A208368 003</u>	Sep 18, 2020
<u>AB</u>	HETERO LABS LTD V	<u>EQ 30MG BASE</u>	<u>A209403 001</u>	Oct 07, 2020
<u>AB</u>		<u>EQ 60MG BASE</u>	<u>A209403 002</u>	Oct 07, 2020
<u>AB</u>		<u>EQ 90MG BASE</u>	<u>A209403 003</u>	Oct 07, 2020
<u>AB</u>	SLATE RUN PHARMA	<u>EQ 30MG BASE</u>	<u>A210207 001</u>	Aug 01, 2018
<u>AB</u>		<u>EQ 60MG BASE</u>	<u>A210207 002</u>	Aug 01, 2018
<u>AB</u>		<u>EQ 90MG BASE</u>	<u>A210207 003</u>	Aug 01, 2018
<u>AB</u>	STRIDES PHARMA	<u>EQ 30MG BASE</u>	<u>A209226 001</u>	Apr 30, 2018
<u>AB</u>		<u>EQ 60MG BASE</u>	<u>A209226 002</u>	Apr 30, 2018
<u>AB</u>	!	<u>EQ 90MG BASE</u>	<u>A209226 003</u>	Apr 30, 2018
<u>AB</u>	SUN PHARM	<u>EQ 60MG BASE</u>	<u>A207008 002</u>	Oct 11, 2018
<u>AB</u>	WATSON LABS TEVA	<u>EQ 30MG BASE</u>	<u>A204377 001</u>	Dec 27, 2018
<u>AB</u>		<u>EQ 60MG BASE</u>	<u>A204377 002</u>	Dec 27, 2018
<u>AB</u>		<u>EQ 90MG BASE</u>	<u>A204377 003</u>	Dec 27, 2018

CIPROFLOXACIN

FOR SUSPENSION; ORAL

CIPRO

<u>AB</u>	+	BAYER HLTHCARE	<u>250MG/5ML</u>	<u>N020780 001</u>	Sep 26, 1997
<u>AB</u>	+	!	<u>500MG/5ML</u>	<u>N020780 002</u>	Sep 26, 1997

CIPROFLOXACIN

<u>AB</u>		CHARTWELL	<u>250MG/5ML</u>	<u>A200563 001</u>	Mar 05, 2014
<u>AB</u>			<u>500MG/5ML</u>	<u>A200563 002</u>	Mar 05, 2014

INJECTABLE; INJECTION

CIPROFLOXACIN

<u>AP</u>	!	BAXTER HLTHCARE CORP	<u>200MG/20ML (10MG/ML)</u>	<u>A078062 001</u>	Apr 29, 2008
-----------	---	----------------------	-----------------------------	--------------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

CIPROFLOXACIN

INJECTABLE; INJECTION

CIPROFLOXACIN

<u>AP</u>	!		<u>400MG/40ML (10MG/ML)</u>	<u>A078062</u>	<u>002</u>	Apr 29, 2008
<u>AP</u>		HIKMA FARMACEUTICA	<u>200MG/20ML (10MG/ML)</u>	<u>A076717</u>	<u>001</u>	Dec 22, 2009
<u>AP</u>			<u>400MG/40ML (10MG/ML)</u>	<u>A076717</u>	<u>002</u>	Dec 22, 2009

CIPROFLOXACIN IN DEXTROSE 5% IN PLASTIC CONTAINER

<u>AP</u>		HIKMA FARMACEUTICA	<u>400MG/200ML</u>	<u>A078431</u>	<u>001</u>	Nov 18, 2009
<u>AP</u>	!	HOSPIRA	<u>200MG/100ML</u>	<u>A077753</u>	<u>001</u>	Mar 18, 2008
<u>AP</u>	!		<u>400MG/200ML</u>	<u>A077753</u>	<u>002</u>	Mar 18, 2008
<u>AP</u>		INFORLIFE	<u>200MG/100ML</u>	<u>A078252</u>	<u>001</u>	Mar 18, 2008
<u>AP</u>			<u>400MG/200ML</u>	<u>A078252</u>	<u>002</u>	Mar 18, 2008

CIPROFLOXACIN HYDROCHLORIDE

OINTMENT; OPHTHALMIC

CILOXAN

+! SANDOZ

EQ 0.3% BASE

N020369 001 Mar 30, 1998

SOLUTION/DROPS; OPHTHALMIC

CILOXAN

<u>AT</u>	+!	SANDOZ	<u>EQ 0.3% BASE</u>	<u>N019992</u>	<u>001</u>	Dec 31, 1990
-----------	----	--------	---------------------	----------------	------------	--------------

CIPROFLOXACIN HYDROCHLORIDE

<u>AT</u>		ALTAIRE PHARMS INC	<u>EQ 0.3% BASE</u>	<u>A204613</u>	<u>001</u>	May 03, 2018
<u>AT</u>		FDC LTD	<u>EQ 0.3% BASE</u>	<u>A077568</u>	<u>001</u>	Jun 30, 2008
<u>AT</u>		RISING	<u>EQ 0.3% BASE</u>	<u>A077689</u>	<u>001</u>	Dec 13, 2006
<u>AT</u>		RUBICON	<u>EQ 0.3% BASE</u>	<u>A075928</u>	<u>001</u>	Jun 09, 2004
<u>AT</u>		WATSON LABS INC	<u>EQ 0.3% BASE</u>	<u>A076673</u>	<u>001</u>	Jan 21, 2005

SOLUTION/DROPS; OTIC

CETRAHAL

<u>AB</u>	+!	KEY THERAP	<u>EQ 0.2% BASE</u>	<u>N021918</u>	<u>001</u>	May 01, 2009
-----------	----	------------	---------------------	----------------	------------	--------------

CIPROFLOXACIN HYDROCHLORIDE

<u>AB</u>		IDENTIRX	<u>EQ 0.2% BASE</u>	<u>A217887</u>	<u>001</u>	Nov 05, 2024
-----------	--	----------	---------------------	----------------	------------	--------------

TABLET; ORAL

CIPRO

<u>AB</u>	+	BAYER HLTHCARE	<u>EQ 250MG BASE</u>	<u>N019537</u>	<u>002</u>	Oct 22, 1987
-----------	---	----------------	----------------------	----------------	------------	--------------

<u>AB</u>	+!		<u>EQ 500MG BASE</u>	<u>N019537</u>	<u>003</u>	Oct 22, 1987
-----------	----	--	----------------------	----------------	------------	--------------

CIPROFLOXACIN HYDROCHLORIDE

<u>AB</u>		AIPING PHARM INC	<u>EQ 250MG BASE</u>	<u>A076593</u>	<u>002</u>	Jun 09, 2004
<u>AB</u>			<u>EQ 500MG BASE</u>	<u>A076593</u>	<u>003</u>	Jun 09, 2004
<u>AB</u>			<u>EQ 750MG BASE</u>	<u>A076593</u>	<u>004</u>	Jun 09, 2004
<u>AB</u>		AMNEAL	<u>EQ 250MG BASE</u>	<u>A075939</u>	<u>002</u>	Jun 09, 2004
<u>AB</u>			<u>EQ 500MG BASE</u>	<u>A075939</u>	<u>003</u>	Jun 09, 2004
<u>AB</u>			<u>EQ 750MG BASE</u>	<u>A075939</u>	<u>004</u>	Jun 09, 2004
<u>AB</u>		AUROBINDO PHARMA	<u>EQ 250MG BASE</u>	<u>A077859</u>	<u>001</u>	Apr 26, 2007
<u>AB</u>			<u>EQ 500MG BASE</u>	<u>A077859</u>	<u>002</u>	Apr 26, 2007
<u>AB</u>			<u>EQ 750MG BASE</u>	<u>A077859</u>	<u>003</u>	Apr 26, 2007
<u>AB</u>		CARLSBAD	<u>EQ 250MG BASE</u>	<u>A076126</u>	<u>002</u>	Jun 09, 2004
<u>AB</u>			<u>EQ 500MG BASE</u>	<u>A076126</u>	<u>003</u>	Jun 09, 2004
<u>AB</u>			<u>EQ 750MG BASE</u>	<u>A076126</u>	<u>004</u>	Jun 09, 2004
<u>AB</u>		CHARTWELL	<u>EQ 250MG BASE</u>	<u>A076896</u>	<u>001</u>	Nov 04, 2004
<u>AB</u>			<u>EQ 500MG BASE</u>	<u>A076896</u>	<u>002</u>	Nov 04, 2004
<u>AB</u>			<u>EQ 750MG BASE</u>	<u>A076896</u>	<u>003</u>	Nov 04, 2004
<u>AB</u>		DR REDDYS LABS LTD	<u>EQ 250MG BASE</u>	<u>A075593</u>	<u>003</u>	Jun 09, 2004
<u>AB</u>			<u>EQ 500MG BASE</u>	<u>A075593</u>	<u>004</u>	Jun 09, 2004
<u>AB</u>			<u>EQ 750MG BASE</u>	<u>A075593</u>	<u>001</u>	Jun 09, 2004
<u>AB</u>		HIKMA	<u>EQ 250MG BASE</u>	<u>A076558</u>	<u>002</u>	Jun 09, 2004
<u>AB</u>			<u>EQ 500MG BASE</u>	<u>A076558</u>	<u>003</u>	Jun 09, 2004
<u>AB</u>			<u>EQ 750MG BASE</u>	<u>A076558</u>	<u>004</u>	Jun 09, 2004
<u>AB</u>		IVAX SUB TEVA PHARMS	<u>EQ 250MG BASE</u>	<u>A076089</u>	<u>002</u>	Jun 09, 2004
<u>AB</u>			<u>EQ 500MG BASE</u>	<u>A076089</u>	<u>003</u>	Jun 09, 2004
<u>AB</u>			<u>EQ 750MG BASE</u>	<u>A076089</u>	<u>004</u>	Jun 09, 2004
<u>AB</u>		RISING	<u>EQ 500MG BASE</u>	<u>A075817</u>	<u>003</u>	Jun 09, 2004
<u>AB</u>		UNIQUE	<u>EQ 250MG BASE</u>	<u>A076639</u>	<u>001</u>	Sep 10, 2004
<u>AB</u>			<u>EQ 500MG BASE</u>	<u>A076639</u>	<u>002</u>	Sep 10, 2004
<u>AB</u>			<u>EQ 750MG BASE</u>	<u>A076639</u>	<u>003</u>	Sep 10, 2004
<u>AB</u>		WATSON LABS	<u>EQ 250MG BASE</u>	<u>A076794</u>	<u>002</u>	Jun 09, 2004
<u>AB</u>			<u>EQ 500MG BASE</u>	<u>A076794</u>	<u>003</u>	Jun 09, 2004
<u>AB</u>			<u>EQ 750MG BASE</u>	<u>A076794</u>	<u>004</u>	Jun 09, 2004
<u>AB</u>		YILING	<u>EQ 250MG BASE</u>	<u>A208921</u>	<u>001</u>	Jun 22, 2018
<u>AB</u>			<u>EQ 500MG BASE</u>	<u>A208921</u>	<u>002</u>	Jun 22, 2018

PRESCRIPTION DRUG PRODUCT LIST

CIPROFLOXACIN HYDROCHLORIDE; FLUOCINOLONE ACETONIDE

SOLUTION/DROPS;OTIC

OTOVEL

+! LABORATORIOS SALVAT EQ 0.3% BASE;0.025%

N208251 001 Apr 29, 2016

CIPROFLOXACIN HYDROCHLORIDE; HYDROCORTISONE

SUSPENSION/DROPS;OTIC

CIPRO HC

+! SANDOZ EQ 0.2% BASE;1%

N020805 001 Feb 10, 1998

CIPROFLOXACIN; CIPROFLOXACIN HYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL

CIPROFLOXACIN EXTENDED RELEASE

! DR REDDYS LABS LTD 425.2MG;EQ 574.9MG BASE

A077701 001 Mar 26, 2007

CIPROFLOXACIN; DEXAMETHASONE

SUSPENSION/DROPS;OTIC

CIPRODEXAB +! SANDOZ 0.3%;0.1%N021537 001 Jul 18, 2003CIPROFLOXACIN AND DEXAMETHASONEAB DR REDDYS 0.3%;0.1%A205548 001 Aug 10, 2020AB SENTISS 0.3%;0.1%A215768 001 Jun 09, 2023AB SUN PHARM 0.3%;0.1%A210470 001 Aug 30, 2022CIPROFLOXACIN; DEXAMETHASONEAB AMNEAL 0.3%;0.1%A216501 001 Mar 22, 2024CISATRACURIUM BESYLATE

INJECTABLE; INJECTION

CISATRACURIUM BESYLATEAP CAPLIN EQ 2MG BASE/MLA217725 001 Jun 09, 2023AP EQ 10MG BASE/MLA217725 002 Jun 09, 2023AP EUGIA PHARMA EQ 2MG BASE/MLA209144 001 May 08, 2020AP FRESENIUS KABI USA EQ 2MG BASE/MLA203183 001 Feb 26, 2015AP HAINAN POLY EQ 2MG BASE/MLA216538 001 Mar 08, 2024AP HENGRUI PHARMA EQ 2MG BASE/MLA209334 001 Aug 30, 2017AP HIKMA EQ 2MG BASE/MLA203078 001 Jun 28, 2022AP EQ 2MG BASE/MLA203079 001 Apr 19, 2023AP EQ 10MG BASE/MLA203078 002 Dec 18, 2023AP HOSPIRA EQ 2MG BASE/MLA203236 001 Mar 30, 2018AP EQ 2MG BASE/MLA203238 001 Mar 30, 2018AP EQ 10MG BASE/MLA203236 002 Mar 30, 2018AP ! MEITHEAL EQ 2MG BASE/MLA211668 001 Apr 25, 2019AP ! EQ 2MG BASE/MLA211669 001 Apr 25, 2019AP ! EQ 10MG BASE/MLA211668 002 Apr 25, 2019AP PIRAMAL CRITICAL EQ 2MG BASE/MLA215516 001 May 24, 2022AP EQ 2MG BASE/MLA215516 003 Dec 14, 2022AP EQ 10MG BASE/MLA215516 002 May 24, 2022AP SAGENT PHARMS INC EQ 2MG BASE/MLA201836 001 Nov 06, 2020AP EQ 2MG BASE/MLA201851 001 Nov 06, 2020AP EQ 10MG BASE/MLA201836 002 Nov 06, 2020AP SANDOZ INC EQ 2MG BASE/MLA200159 001 Feb 03, 2012AP SOMERSET EQ 2MG BASE/MLA209132 001 Apr 24, 2019AP ZYDUS PHARMS EQ 2MG BASE/MLA212171 001 Nov 04, 2019AP EQ 10MG BASE/MLA212171 002 Nov 04, 2019CISATRACURIUM BESYLATE PRESERVATIVE FREEAP EUGIA PHARMA EQ 2MG BASE/MLA209665 001 Oct 27, 2020AP FRESENIUS KABI USA EQ 2MG BASE/MLA203182 001 Feb 26, 2015AP EQ 10MG BASE/MLA203182 002 Feb 26, 2015AP HAINAN POLY EQ 2MG BASE/MLA216539 001 Aug 27, 2024AP EQ 10MG BASE/MLA216539 002 Aug 27, 2024AP HENGRUI PHARMA EQ 2MG BASE/MLA204960 001 Jan 27, 2017AP EQ 10MG BASE/MLA204960 002 Sep 19, 2017AP SANDOZ INC EQ 2MG BASE/MLA200154 001 Feb 03, 2012AP EQ 10MG BASE/MLA200154 002 Feb 03, 2012AP SOMERSET THERAPS EQ 2MG BASE/MLA206791 001 Feb 20, 2019AP LLC EQ 10MG BASE/MLA206791 002 Feb 20, 2019CISPLATIN

INJECTABLE; INJECTION

CISPLATINAP ACCORD HLTHCARE 1MG/MLA206774 001 Aug 18, 2015AP ! FRESENIUS KABI USA 1MG/MLA074735 001 Jul 16, 1999AP GLAND PHARMA LTD 1MG/MLA207323 001 Mar 17, 2017AP HIKMA 1MG/MLA075036 001 Nov 07, 2000

PRESCRIPTION DRUG PRODUCT LIST

CISPLATIN

INJECTABLE; INJECTION

CISPLATIN

<u>AP</u>	+	HQ SPCLT PHARMA	<u>1MG/ML</u>	<u>N018057 004</u>	Nov 08, 1988
<u>AP</u>		PHARMACHEMIE BV	<u>1MG/ML</u>	<u>A074656 001</u>	May 16, 2000
<u>AP</u>		QILU	<u>1MG/ML</u>	<u>A218868 001</u>	May 10, 2024
	+	HQ SPCLT PHARMA	50MG/VIAL	N018057 002	

CITALOPRAM HYDROBROMIDE

CAPSULE; ORAL

CITALOPRAM HYDROBROMIDE

+! ALMATICA

EQ 30MG BASE

N215428 001 Jan 31, 2022

SOLUTION; ORAL

CITALOPRAM HYDROBROMIDE

<u>AA</u>		AUROBINDO PHARMA	<u>EQ 10MG BASE/5ML</u>	<u>A077812 001</u>	Aug 28, 2006
<u>AA</u>		CHARTWELL MOLECULAR	<u>EQ 10MG BASE/5ML</u>	<u>A077629 001</u>	Jun 14, 2006
<u>AA</u>		HETERO LABS LTD III	<u>EQ 10MG BASE/5ML</u>	<u>A201450 001</u>	Dec 15, 2015
<u>AA</u>	!	HIKMA	<u>EQ 10MG BASE/5ML</u>	<u>A077043 001</u>	Dec 13, 2004

TABLET; ORAL

CELEXA

<u>AB</u>	+	ABEVIE	<u>EQ 10MG BASE</u>	<u>N020822 001</u>	Apr 27, 2000
<u>AB</u>	+		<u>EQ 20MG BASE</u>	<u>N020822 002</u>	Jul 17, 1998
<u>AB</u>	+		<u>EQ 40MG BASE</u>	<u>N020822 003</u>	Jul 17, 1998

CITALOPRAM HYDROBROMIDE

<u>AB</u>		AMNEAL PHARMS NY	<u>EQ 10MG BASE</u>	<u>A077289 001</u>	Nov 30, 2006
<u>AB</u>			<u>EQ 20MG BASE</u>	<u>A077289 002</u>	Nov 30, 2006
<u>AB</u>			<u>EQ 40MG BASE</u>	<u>A077289 003</u>	Nov 30, 2006
<u>AB</u>		APOTEX	<u>EQ 10MG BASE</u>	<u>A077046 001</u>	Nov 24, 2004
<u>AB</u>			<u>EQ 20MG BASE</u>	<u>A077046 002</u>	Nov 24, 2004
<u>AB</u>			<u>EQ 40MG BASE</u>	<u>A077046 003</u>	Nov 24, 2004
<u>AB</u>		AUROBINDO	<u>EQ 10MG BASE</u>	<u>A077031 001</u>	Oct 28, 2004
<u>AB</u>			<u>EQ 20MG BASE</u>	<u>A077031 002</u>	Oct 28, 2004
<u>AB</u>			<u>EQ 40MG BASE</u>	<u>A077031 003</u>	Oct 28, 2004
<u>AB</u>		CHARTWELL MOLECULAR	<u>EQ 10MG BASE</u>	<u>A077044 001</u>	Nov 05, 2004
<u>AB</u>			<u>EQ 20MG BASE</u>	<u>A077044 002</u>	Nov 05, 2004
<u>AB</u>			<u>EQ 40MG BASE</u>	<u>A077044 003</u>	Nov 05, 2004
<u>AB</u>		DR REDDYS LABS LTD	<u>EQ 10MG BASE</u>	<u>A077038 001</u>	Oct 28, 2004
<u>AB</u>			<u>EQ 20MG BASE</u>	<u>A077038 002</u>	Oct 28, 2004
<u>AB</u>			<u>EQ 40MG BASE</u>	<u>A077038 003</u>	Oct 28, 2004
<u>AB</u>		EPIC PHARMA	<u>EQ 10MG BASE</u>	<u>A077045 003</u>	Apr 29, 2005
<u>AB</u>			<u>EQ 20MG BASE</u>	<u>A077045 002</u>	Apr 29, 2005
<u>AB</u>			<u>EQ 40MG BASE</u>	<u>A077045 001</u>	Apr 29, 2005
<u>AB</u>		GLENMARK PHARMS LTD	<u>EQ 10MG BASE</u>	<u>A077654 001</u>	Feb 27, 2009
<u>AB</u>			<u>EQ 20MG BASE</u>	<u>A077654 002</u>	Feb 27, 2009
<u>AB</u>			<u>EQ 40MG BASE</u>	<u>A077654 003</u>	Feb 27, 2009
<u>AB</u>		INVAGEN PHARMS	<u>EQ 10MG BASE</u>	<u>A077534 001</u>	Oct 03, 2006
<u>AB</u>			<u>EQ 20MG BASE</u>	<u>A077534 002</u>	Oct 03, 2006
<u>AB</u>			<u>EQ 40MG BASE</u>	<u>A077534 003</u>	Oct 03, 2006
<u>AB</u>		MYLAN	<u>EQ 10MG BASE</u>	<u>A077042 001</u>	Nov 05, 2004
<u>AB</u>			<u>EQ 20MG BASE</u>	<u>A077042 002</u>	Nov 05, 2004
<u>AB</u>			<u>EQ 40MG BASE</u>	<u>A077042 003</u>	Nov 05, 2004
<u>AB</u>		TORRENT PHARMS	<u>EQ 10MG BASE</u>	<u>A078216 001</u>	Mar 27, 2007
<u>AB</u>			<u>EQ 20MG BASE</u>	<u>A078216 002</u>	Mar 27, 2007
<u>AB</u>			<u>EQ 40MG BASE</u>	<u>A078216 003</u>	Mar 27, 2007

CITRIC ACID; GLUCONOLACTONE; MAGNESIUM CARBONATE

SOLUTION; IRRIGATION

RENACIDIN

+! UNITED GUARDIAN

6.602GM/100ML;198MG/100ML;3.177GM/100ML

N019481 001 Oct 02, 1990

CITRIC ACID; LACTIC ACID; POTASSIUM BITARTRATE

GEL; VAGINAL

PHEXXI

+! EVOFEM INC

1%;1.8%;0.4%

N208352 001 May 22, 2020

CITRIC ACID; MAGNESIUM OXIDE; SODIUM PICOSULFATE

FOR SOLUTION; ORAL

SODIUM PICOSULFATE, MAGNESIUM OXIDE AND ANHYDROUS CITRIC ACID

! HETERO LABS LTD V

12GM/PACKET;3.5GM/PACKET;10MG/PACKET

A212789 001 Jul 18, 2022

SOLUTION; ORAL

CLENPIQ

+! FERRING PHARMS INC

12GM/BOT;3.5GM/BOT;10MG/BOT

N209589 001 Nov 28, 2017

PRESCRIPTION DRUG PRODUCT LIST

CITRIC ACID; UREA C-13

FOR SOLUTION, TABLET, FOR SOLUTION;ORAL

IDKIT:HP

+! MERIDIAN BIOSCIENCE 4GM;75MG

N021314 001 Dec 17, 2002

CLADRIBINE

INJECTABLE; INJECTION

CLADRIBINE**AP** ! FRESENIUS KABI USA1MG/ML**A076571 001** Apr 22, 2004**AP** HIKMA1MG/ML**A075405 001** Feb 28, 2000**AP** HISUN PHARM
HANGZHOU1MG/ML**A210856 001** Nov 25, 2019

TABLET;ORAL

MAVENCLAD

+! EMD SERONO INC

10MG

N022561 001 Mar 29, 2019

CLARITHROMYCIN

FOR SUSPENSION;ORAL

CLARITHROMYCIN

SANDOZ

125MG/5ML

A065283 002 Sep 04, 2007

!

250MG/5ML

A065283 003 Sep 04, 2007

TABLET;ORAL

CLARITHROMYCIN**AB** ALEMBIC250MG**A210459 001** Jan 31, 2022**AB**500MG**A210459 002** Jan 31, 2022**AB** ! AUROBINDO250MG**A065489 001** Jul 25, 2012**AB** !500MG**A065489 002** Jul 25, 2012**AB** CHARTWELL250MG**A065384 001** Aug 20, 2007**AB**500MG**A065384 002** Aug 20, 2007**AB** ESJAY PHARMA250MG**A202710 001** Jun 10, 2013**AB**500MG**A202710 002** Jun 10, 2013**AB** HEC PHARM250MG**A203584 001** Sep 28, 2015**AB**500MG**A203584 002** Sep 28, 2015**AB** SANDOZ250MG**A065144 001** Oct 18, 2005**AB**500MG**A065136 001** Aug 25, 2005

TABLET, EXTENDED RELEASE;ORAL

CLARITHROMYCIN**AB** ACTAVIS LABS FL INC500MG**A065145 001** Jun 24, 2004**AB** ! DR REDDYS LABS SA500MG**A065154 001** May 18, 2005**AB** NOSTRUM LABS INC500MG**A203243 001** Feb 29, 2016**AB** SUNSHINE500MG**A208987 001** Jul 09, 2018CLASCOTERONE

CREAM;TOPICAL

WINLEVI

+! SUN PHARM

1%

N213433 001 Aug 26, 2020

CLEMASTINE FUMARATE

SYRUP;ORAL

CLEMASTINE FUMARATE**AA** ! GENUSEQ 0.5MG BASE/5ML**A073399 001** Jun 30, 1994**AA** NEW HEIGHTSRXEQ 0.5MG BASE/5ML**A074884 001** Dec 17, 1997

TABLET;ORAL

CLEMASTINE FUMARATE

! GENUS

2.68MG

A073283 001 Jan 31, 1992

CLEVIDIPINE

EMULSION; INTRAVENOUS

CLEVIPREX

+! CHIESI

25MG/50ML (0.5MG/ML)

N022156 001 Aug 01, 2008

+!

50MG/100ML (0.5MG/ML)

N022156 002 Aug 01, 2008

CLINDAMYCIN HYDROCHLORIDE

CAPSULE;ORAL

CLEOCIN HYDROCHLORIDE**AB** + PFIZEREQ 75MG BASE**N050162 001****AB** +EQ 150MG BASE**N050162 002****AB** +!EQ 300MG BASE**N050162 003** Apr 14, 1988CLINDAMYCIN HYDROCHLORIDE**AB** AUROBINDO PHARMAEQ 150MG BASE**A065442 001** Aug 26, 2009**AB**EQ 300MG BASE**A065442 002** Aug 26, 2009**AB** CHARTWELL MOLECULAREQ 75MG BASE**A065243 002** Aug 12, 2005**AB** EPIC PHARMA LLCEQ 150MG BASE**A065194 001** Mar 22, 2004**AB**EQ 300MG BASE**A065194 002** Mar 22, 2004**AB** GLENMARK PHARMS LTDEQ 75MG BASE**A216957 001** Mar 10, 2023**AB**EQ 150MG BASE**A216957 002** Mar 10, 2023

PRESCRIPTION DRUG PRODUCT LIST

CLINDAMYCIN HYDROCHLORIDE

CAPSULE; ORAL

CLINDAMYCIN HYDROCHLORIDE

<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A216957 003</u>	Mar 10, 2023
<u>AB</u>	MICRO LABS	<u>EQ 75MG BASE</u>	<u>A207402 001</u>	Nov 05, 2018
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A207402 002</u>	Nov 05, 2018
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A207402 003</u>	Nov 05, 2018
<u>AB</u>	SUN PHARM INDS LTD	<u>EQ 150MG BASE</u>	<u>A065061 001</u>	Feb 02, 2001
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A065061 002</u>	Feb 02, 2001
<u>AB</u>	ZYDUS PHARMS USA	<u>EQ 75MG BASE</u>	<u>A065217 001</u>	Jan 31, 2005
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A065217 002</u>	Jan 31, 2005
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A065217 003</u>	Jan 31, 2005

CLINDAMYCIN PALMITATE HYDROCHLORIDE

FOR SOLUTION; ORAL

CLEOCIN

<u>AA</u>	!	PFIZER	<u>EQ 75MG BASE/5ML</u>	<u>A062644 001</u>	Apr 07, 1986
-----------	---	--------	-------------------------	--------------------	--------------

CLINDAMYCIN PALMITATE HYDROCHLORIDE

<u>AA</u>		AMNEAL PHARMS	<u>EQ 75MG BASE/5ML</u>	<u>A203513 001</u>	Mar 13, 2014
<u>AA</u>		APPCO	<u>EQ 75MG BASE/5ML</u>	<u>A207047 001</u>	May 11, 2018
<u>AA</u>		AUROBINDO PHARMA LTD	<u>EQ 75MG BASE/5ML</u>	<u>A202409 001</u>	Apr 30, 2013
<u>AA</u>		CHARTWELL RX	<u>EQ 75MG BASE/5ML</u>	<u>A206958 001</u>	May 05, 2017
<u>AA</u>		PADAGIS US	<u>EQ 75MG BASE/5ML</u>	<u>A090902 001</u>	Jul 07, 2010
<u>AA</u>		RISING	<u>EQ 75MG BASE/5ML</u>	<u>A201821 001</u>	Aug 28, 2012

CLINDAMYCIN PHOSPHATE

AEROSOL, FOAM; TOPICAL

CLINDAMYCIN PHOSPHATE

<u>AT</u>		GLENMARK PHARMS LTD	<u>1%</u>	<u>A210778 001</u>	Sep 20, 2021
<u>AT</u>	!	PADAGIS ISRAEL	<u>1%</u>	<u>A090785 001</u>	Mar 31, 2010
<u>AT</u>		TARO	<u>1%</u>	<u>A210004 001</u>	Mar 11, 2020

CREAM; VAGINAL

CLEOCIN

<u>AB</u>	+	!	PFIZER	<u>EQ 2% BASE</u>	<u>N050680 002</u>	Mar 02, 1998
-----------	---	---	--------	-------------------	--------------------	--------------

CLINDAMYCIN PHOSPHATE

<u>AB</u>		FOUGERA PHARMS	<u>EQ 2% BASE</u>	<u>A065139 001</u>	Dec 27, 2004	
		CLINDESSE				
	+	!	PADAGIS US	<u>EQ 2% BASE</u>	<u>N050793 001</u>	Nov 30, 2004

GEL; TOPICAL

CLEOCIN T

<u>AB1</u>	+	!	PFIZER	<u>EQ 1% BASE</u>	<u>N050615 001</u>	Jan 07, 1987
------------	---	---	--------	-------------------	--------------------	--------------

CLINDAMYCIN PHOSPHATE

<u>AB1</u>		ALEMBIC	<u>EQ 1% BASE</u>	<u>A212466 001</u>	Apr 02, 2024
<u>AB1</u>		ENCUBE	<u>EQ 1% BASE</u>	<u>A212438 001</u>	Mar 11, 2021
<u>AB1</u>		FOUGERA PHARMS	<u>EQ 1% BASE</u>	<u>A064160 001</u>	Jan 28, 2000
<u>AB1</u>		GLENMARK PHARMS LTD	<u>EQ 1% BASE</u>	<u>A214251 001</u>	Feb 10, 2021
<u>AB1</u>		PADAGIS ISRAEL	<u>EQ 1% BASE</u>	<u>A212104 001</u>	Dec 31, 2020
<u>AB1</u>		QUAGEN	<u>EQ 1% BASE</u>	<u>A211872 001</u>	Jul 29, 2020
<u>AB1</u>		TARO	<u>EQ 1% BASE</u>	<u>A214052 001</u>	Nov 10, 2020
<u>AB1</u>		ZYDUS LIFESCIENCES	<u>EQ 1% BASE</u>	<u>A216587 001</u>	Sep 19, 2023

CLINDAGEL

<u>AB2</u>	+	!	BAUSCH	<u>EQ 1% BASE</u>	<u>N050782 001</u>	Nov 27, 2000
------------	---	---	--------	-------------------	--------------------	--------------

CLINDAMYCIN PHOSPHATE

<u>AB2</u>		AMNEAL	<u>EQ 1% BASE</u>	<u>A214668 001</u>	Aug 05, 2022
<u>AB2</u>		SOLARIS PHARMA CORP	<u>EQ 1% BASE</u>	<u>A212842 001</u>	Aug 13, 2021

GEL; VAGINAL

XACIATO

	+	!	ORGANON LLC	<u>EQ 2% BASE</u>	<u>N215650 001</u>	Dec 07, 2021
--	---	---	-------------	-------------------	--------------------	--------------

INJECTABLE; INJECTION

CLEOCIN PHOSPHATE

<u>AP</u>		PFIZER	<u>EQ 150MG BASE/ML</u>	<u>A062803 001</u>	Oct 16, 1987
<u>AP</u>	+	!		<u>EQ 150MG BASE/ML</u>	<u>N050441 001</u>

CLINDAMYCIN PHOSPHATE

<u>AP</u>		ALMAJECT	<u>EQ 150MG BASE/ML</u>	<u>A062800 001</u>	Jul 24, 1987
<u>AP</u>			<u>EQ 150MG BASE/ML</u>	<u>A062943 001</u>	Sep 29, 1988
<u>AP</u>		FRESENIUS KABI USA	<u>EQ 150MG BASE/ML</u>	<u>A065346 001</u>	Mar 29, 2007
<u>AP</u>			<u>EQ 150MG BASE/ML</u>	<u>A065347 001</u>	May 09, 2007
<u>AP</u>		HIKMA	<u>EQ 150MG BASE/ML</u>	<u>A062889 001</u>	Apr 25, 1988
<u>AP</u>			<u>EQ 150MG BASE/ML</u>	<u>A065206 001</u>	Sep 24, 2004
<u>AP</u>		SAGENT PHARMS INC	<u>EQ 150MG BASE/ML</u>	<u>A090108 001</u>	Sep 30, 2011
<u>AP</u>			<u>EQ 150MG BASE/ML</u>	<u>A090109 001</u>	Sep 30, 2011

PRESCRIPTION DRUG PRODUCT LIST

CLINDAMYCIN PHOSPHATE

INJECTABLE; INJECTION

CLINDAMYCIN PHOSPHATE IN 5% DEXTROSE IN PLASTIC CONTAINER

<u>AP</u>	BAXTER HLTHCARE CORP	<u>EQ 6MG BASE/ML</u>	<u>A208084 001</u>	Jun 28, 2017
<u>AP</u>		<u>EQ 12MG BASE/ML</u>	<u>A208084 002</u>	Jun 28, 2017
<u>AP</u>		<u>EQ 18MG BASE/ML</u>	<u>A208084 003</u>	Jun 28, 2017
<u>AP</u>	RISING	<u>EQ 6MG BASE/ML</u>	<u>A203048 001</u>	Apr 04, 2013
<u>AP</u>		<u>EQ 12MG BASE/ML</u>	<u>A203048 002</u>	Apr 04, 2013
<u>AP</u>		<u>EQ 18MG BASE/ML</u>	<u>A203048 003</u>	Apr 04, 2013
<u>AP</u>	! SANDOZ INC	<u>EQ 6MG BASE/ML</u>	<u>A201692 001</u>	May 31, 2012
<u>AP</u>	!	<u>EQ 12MG BASE/ML</u>	<u>A201692 002</u>	May 31, 2012
<u>AP</u>	!	<u>EQ 18MG BASE/ML</u>	<u>A201692 003</u>	May 31, 2012

CLINDAMYCIN PHOSPHATE IN DEXTROSE 5%

<u>AP</u>	HIKMA	<u>EQ 6MG BASE/ML</u>	<u>A214401 001</u>	Apr 05, 2023
<u>AP</u>		<u>EQ 12MG BASE/ML</u>	<u>A214401 002</u>	Apr 05, 2023
<u>AP</u>		<u>EQ 18MG BASE/ML</u>	<u>A214401 003</u>	Apr 05, 2023
	+! ABRAXIS PHARM	EQ 900MG BASE/100ML	N050635 001	Dec 22, 1989

LOTION; TOPICAL

CLEOCIN T

<u>AB</u>	+! PFIZER	<u>EQ 1% BASE</u>	<u>N050600 001</u>	May 31, 1989
-----------	-----------	-------------------	--------------------	--------------

CLINDAMYCIN PHOSPHATE

<u>AB</u>	ENCUBE	<u>EQ 1% BASE</u>	<u>A215607 001</u>	Jan 18, 2022
<u>AB</u>	FOUGERA PHARMS	<u>EQ 1% BASE</u>	<u>A065067 001</u>	Jan 31, 2002
<u>AB</u>	PADAGIS ISRAEL	<u>EQ 1% BASE</u>	<u>A214604 001</u>	Mar 08, 2021
<u>AB</u>	TARO	<u>EQ 1% BASE</u>	<u>A214526 001</u>	Jun 28, 2021

SOLUTION; INTRAVENOUS

CLINDAMYCIN PHOSPHATE IN 0.9% SODIUM CHLORIDE

	+! BAXTER HLTHCARE CORP	EQ 300MG BASE/50ML (EQ 6MG BASE/ML)	N208083 001	Apr 20, 2017
	+!	EQ 600MG BASE/50ML (EQ 12MG BASE/ML)	N208083 002	Apr 20, 2017
	+!	EQ 900MG BASE/50ML (EQ 18MG BASE/ML)	N208083 003	Apr 20, 2017

SOLUTION; TOPICAL

CLINDA-DERM

<u>AT</u>	PADAGIS US	<u>EQ 1% BASE</u>	<u>A063329 001</u>	Sep 30, 1992
-----------	------------	-------------------	--------------------	--------------

CLINDAMYCIN PHOSPHATE

<u>AT</u>	CHARTWELL RX	<u>EQ 1% BASE</u>	<u>A209846 001</u>	Feb 08, 2018
<u>AT</u>	ENCUBE ETHICALS	<u>EQ 1% BASE</u>	<u>A209914 001</u>	Jan 28, 2019
<u>AT</u>	FOUGERA PHARMS INC	<u>EQ 1% BASE</u>	<u>A064159 001</u>	Jun 05, 1997
<u>AT</u>	! PADAGIS US	<u>EQ 1% BASE</u>	<u>A064050 001</u>	Nov 30, 1995
<u>AT</u>	TARO PHARM INDS	<u>EQ 1% BASE</u>	<u>A065184 001</u>	Mar 31, 2004
<u>AT</u>	ZYDUS LIFESCIENCES	<u>EQ 1% BASE</u>	<u>A208767 001</u>	Jul 16, 2018

SUPPOSITORY; VAGINAL

CLEOCIN

	+! PFIZER	100MG	N050767 001	Aug 13, 1999
--	-----------	-------	-------------	--------------

SWAB; TOPICAL

CLINDAMYCIN PHOSPHATE

<u>AT</u>	EPIC PHARMA LLC	<u>EQ 1% BASE</u>	<u>A065513 001</u>	Jun 17, 2010
<u>AT</u>	! PADAGIS US	<u>EQ 1% BASE</u>	<u>A065049 001</u>	May 25, 2000

CLINDETS

<u>AT</u>	PADAGIS US	<u>EQ 1% BASE</u>	<u>A064136 001</u>	Sep 30, 1996
-----------	------------	-------------------	--------------------	--------------

CLINDAMYCIN PHOSPHATE; TRETINOIN

GEL; TOPICAL

CLINDAMYCIN PHOSPHATE AND TRETINOIN

<u>AB1</u>	ACTAVIS MID ATLANTIC	<u>1.2%;0.025%</u>	<u>A202564 001</u>	Jun 12, 2015
------------	-------------------------	--------------------	--------------------	--------------

ZIANA

<u>AB1</u>	+! BAUSCH	<u>1.2%;0.025%</u>	<u>N050802 001</u>	Nov 07, 2006
------------	-----------	--------------------	--------------------	--------------

CLINDAMYCIN PHOSPHATE AND TRETINOIN

<u>AB2</u>	ENCUBE	<u>1.2%;0.025%</u>	<u>A216943 001</u>	Sep 01, 2023
<u>AB2</u>	SOLARIS PHARMA CORP	<u>1.2%;0.025%</u>	<u>A212845 001</u>	Feb 10, 2022

VELTIN

<u>AB2</u>	+! ALMIRALL	<u>1.2%;0.025%</u>	<u>N050803 001</u>	Jul 16, 2010
------------	-------------	--------------------	--------------------	--------------

CLOBAZAM

FILM; ORAL

SYMPAZAN

	+ OTTER PHARMS	5MG	N210833 001	Nov 01, 2018
	+	10MG	N210833 002	Nov 01, 2018
	+!	20MG	N210833 003	Nov 01, 2018

PRESCRIPTION DRUG PRODUCT LIST

CLOBAZAM

SUSPENSION; ORAL

CLOBAZAM

AB	ALKEM LABS LTD	<u>2.5MG/ML</u>	<u>A213039</u>	<u>001</u>	May 06, 2021
AB	AMNEAL	<u>2.5MG/ML</u>	<u>A210039</u>	<u>001</u>	Oct 22, 2018
AB	AUROBINDO PHARMA LTD	<u>2.5MG/ML</u>	<u>A214404</u>	<u>001</u>	Mar 24, 2022
AB	BIONPHARMA	<u>2.5MG/ML</u>	<u>A208819</u>	<u>001</u>	Oct 22, 2018
AB	CHARTWELL MOLECULAR	<u>2.5MG/ML</u>	<u>A213110</u>	<u>001</u>	Apr 24, 2020
AB	HETERO LABS LTD III	<u>2.5MG/ML</u>	<u>A209796</u>	<u>001</u>	Feb 24, 2020
AB	LUPIN LTD	<u>2.5MG/ML</u>	<u>A210546</u>	<u>001</u>	Dec 28, 2018
AB	TARO	<u>2.5MG/ML</u>	<u>A210978</u>	<u>001</u>	Apr 15, 2019

ONFI

AB	+ !	LUNDBECK PHARMS LLC	<u>2.5MG/ML</u>	<u>N203993</u>	<u>001</u>	Dec 14, 2012
-----------	------------	---------------------	-----------------	----------------	------------	--------------

TABLET; ORAL

CLOBAZAM

AB	ALKEM LABS LTD	<u>10MG</u>	<u>A212714</u>	<u>001</u>	Sep 06, 2019
AB		<u>20MG</u>	<u>A212714</u>	<u>002</u>	Sep 06, 2019
AB	ALLIED	<u>10MG</u>	<u>A209308</u>	<u>001</u>	Oct 22, 2018
AB	AMNEAL PHARMS CO	<u>10MG</u>	<u>A209718</u>	<u>001</u>	Oct 22, 2018
AB		<u>20MG</u>	<u>A209718</u>	<u>002</u>	Oct 22, 2018
AB	BIONPHARMA	<u>10MG</u>	<u>A208825</u>	<u>001</u>	Oct 22, 2018
AB		<u>20MG</u>	<u>A208825</u>	<u>002</u>	Oct 22, 2018
AB	HETERO LABS LTD III	<u>10MG</u>	<u>A209795</u>	<u>001</u>	Oct 22, 2018
AB		<u>20MG</u>	<u>A209795</u>	<u>002</u>	Oct 22, 2018
AB	LUPIN LTD	<u>10MG</u>	<u>A210545</u>	<u>001</u>	Dec 14, 2018
AB		<u>20MG</u>	<u>A210545</u>	<u>002</u>	Dec 14, 2018
AB	MICRO LABS	<u>10MG</u>	<u>A211711</u>	<u>001</u>	Jan 30, 2019
AB		<u>20MG</u>	<u>A211711</u>	<u>002</u>	Jan 30, 2019
AB	MSN	<u>10MG</u>	<u>A213404</u>	<u>001</u>	May 11, 2021
AB		<u>20MG</u>	<u>A213404</u>	<u>002</u>	May 11, 2021
AB	PIRAMAL	<u>10MG</u>	<u>A209808</u>	<u>001</u>	Oct 22, 2018
AB		<u>20MG</u>	<u>A209808</u>	<u>002</u>	Oct 22, 2018
AB	UPSHER SMITH LABS	<u>10MG</u>	<u>A209687</u>	<u>001</u>	Oct 22, 2018
AB		<u>20MG</u>	<u>A209687</u>	<u>002</u>	Oct 22, 2018
AB	ZYDUS PHARMS	<u>10MG</u>	<u>A211449</u>	<u>001</u>	Oct 22, 2018
AB		<u>20MG</u>	<u>A211449</u>	<u>002</u>	Oct 22, 2018

ONFI

AB	+	LUNDBECK PHARMS LLC	<u>10MG</u>	<u>N202067</u>	<u>002</u>	Oct 21, 2011
AB	+ !		<u>20MG</u>	<u>N202067</u>	<u>003</u>	Oct 21, 2011

CLOBAZAM

	ALKEM LABS LTD	5MG	A212714	003	Dec 21, 2020
--	----------------	-----	---------	-----	--------------

CLOBETASOL PROPIONATE

AEROSOL, FOAM; TOPICAL

CLOBETASOL PROPIONATE

AB1	ALEMBIC	<u>0.05%</u>	<u>A215838</u>	<u>001</u>	Apr 21, 2022
AB1	GLENMARK PHARMS LTD	<u>0.05%</u>	<u>A210809</u>	<u>001</u>	Feb 15, 2019
AB1	PADAGIS ISRAEL	<u>0.05%</u>	<u>A077763</u>	<u>001</u>	Mar 10, 2008
AB1	! TARO	<u>0.05%</u>	<u>A208779</u>	<u>001</u>	Oct 04, 2018
AB2	GLENMARK PHARMS LTD	<u>0.05%</u>	<u>A211450</u>	<u>001</u>	Sep 09, 2019
AB2	! PADAGIS ISRAEL	<u>0.05%</u>	<u>A201402</u>	<u>001</u>	Aug 14, 2012
AB2	TARO	<u>0.05%</u>	<u>A208563</u>	<u>001</u>	Mar 14, 2024

CREAM; TOPICAL

CLOBETASOL PROPIONATE

AB1	ALEMBIC	<u>0.05%</u>	<u>A213291</u>	<u>001</u>	Jan 27, 2020
AB1	AMNEAL	<u>0.05%</u>	<u>A211256</u>	<u>001</u>	Dec 26, 2018
AB1	AUROBINDO PHARMA USA	<u>0.05%</u>	<u>A075338</u>	<u>001</u>	Feb 09, 2001
AB1	! ENCUBE	<u>0.05%</u>	<u>A212982</u>	<u>001</u>	Aug 28, 2020
AB1	FOUGERA PHARMS INC	<u>0.05%</u>	<u>A074392</u>	<u>001</u>	Sep 30, 1996
AB1	GLENMARK SPECLT	<u>0.05%</u>	<u>A209095</u>	<u>001</u>	May 10, 2018
AB1	LUPIN LTD	<u>0.05%</u>	<u>A210208</u>	<u>001</u>	Jan 30, 2018
AB1	RISING	<u>0.05%</u>	<u>A211401</u>	<u>001</u>	Jan 11, 2019
AB1	TARO	<u>0.05%</u>	<u>A074249</u>	<u>001</u>	Jul 08, 1996
AB1	TORRENT	<u>0.05%</u>	<u>A211836</u>	<u>001</u>	Dec 30, 2019
AB1	XIROMED	<u>0.05%</u>	<u>A210034</u>	<u>001</u>	Jun 15, 2018
AB1	ZYDUS PHARMS	<u>0.05%</u>	<u>A211074</u>	<u>001</u>	Oct 15, 2018

CLOBETASOL PROPIONATE (EMOLLIENT)

AB2	BEACH PRODS	<u>0.05%</u>	<u>A209411</u>	<u>001</u>	Aug 21, 2017
AB2	! FOUGERA PHARMS	<u>0.05%</u>	<u>A075430</u>	<u>001</u>	May 26, 1999
AB2	TARO	<u>0.05%</u>	<u>A075633</u>	<u>001</u>	May 17, 2000

PRESCRIPTION DRUG PRODUCT LIST

CLOBETASOL PROPIONATE

CREAM; TOPICAL

IMPOYZ

+! PRIMUS PHARMS 0.025% N209483 001 Nov 28, 2017

GEL; TOPICAL

CLOBETASOL PROPIONATE**AB** ! FOUGERA PHARMS **0.05%** **A075368 001** Feb 15, 2000**AB** PADAGIS US **0.05%** **A075027 001** Oct 31, 1997**AB** TARO **0.05%** **A075279 001** May 28, 1999

LOTION; TOPICAL

CLOBETASOL PROPIONATE**AB** ACTAVIS MID **0.05%** **A078223 001** Dec 04, 2008

ATLANTIC

AB LUPIN LTD **0.05%** **A209147 001** Sep 22, 2017**AB** TARO **0.05%** **A200302 001** Jul 02, 2012**AB** ZYDUS LIFESCIENCES **0.05%** **A205249 001** Sep 24, 2019CLOBEX**AB** +! GALDERMA LABS LP **0.05%** **N021535 001** Jul 24, 2003

OINTMENT; TOPICAL

CLOBETASOL PROPIONATE**AB** ALEMBIC **0.05%** **A211800 001** Mar 04, 2019**AB** AUROBINDO PHARMA **0.05%** **A075057 001** Aug 12, 1998

USA

AB COSETTE **0.05%** **A074089 001** Feb 16, 1994**AB** ! ENCUBE **0.05%** **A211295 001** Nov 15, 2019**AB** FOUGERA PHARMS **0.05%** **A074407 001** Feb 23, 1996**AB** GLENMARK SPECLT **0.05%** **A208933 001** Mar 20, 2017**AB** MACLEODS PHARMS LTD **0.05%** **A215990 001** Dec 28, 2023**AB** NOVEL LABS INC **0.05%** **A208841 001** May 04, 2018**AB** TARO **0.05%** **A074248 001** Jul 12, 1996**AB** XIROMED **0.05%** **A209701 001** Apr 17, 2018**AB** ZYDUS PHARMS **0.05%** **A210199 001** Oct 27, 2017

SHAMPOO; TOPICAL

CLOBETASOL PROPIONATE**AB** ALEMBIC **0.05%** **A213290 001** May 18, 2020**AB** AMNEAL **0.05%** **A214895 001** Jun 14, 2021**AB** PADAGIS ISRAEL **0.05%** **A090974 001** Aug 09, 2012**AB** TARO **0.05%** **A214867 001** Mar 25, 2021CLOBEX**AB** +! GALDERMA LABS **0.05%** **N021644 001** Feb 05, 2004

SOLUTION; TOPICAL

CLOBETASOL PROPIONATE**AT** ALEMBIC **0.05%** **A212881 001** Oct 21, 2019**AT** ENCUBE **0.05%** **A076977 001** Aug 05, 2005**AT** FOUGERA PHARMS **0.05%** **A075391 001** Feb 08, 1999**AT** GLENMARK PHARMS LTD **0.05%** **A210190 001** Apr 18, 2018**AT** MACLEODS PHARMS LTD **0.05%** **A209361 001** Oct 25, 2017**AT** NOVEL LABS INC **0.05%** **A206075 001** Nov 23, 2015**AT** QUAGEN **0.05%** **A211240 001** Feb 17, 2022**AT** SAPTALIS PHARMS **0.05%** **A211494 001** Oct 02, 2019**AT** ! TARO **0.05%** **A075224 001** Nov 16, 1998**AT** **0.05%** **A075363 001** Dec 29, 2000

SPRAY; TOPICAL

CLOBETASOL PROPIONATE**AT** ALEMBIC **0.05%** **A211191 001** Oct 02, 2019**AT** GLENMARK SPECLT **0.05%** **A209004 001** Mar 26, 2018**AT** LUPIN LTD **0.05%** **A208125 001** Mar 26, 2018**AT** MACLEODS PHARMS LTD **0.05%** **A214345 001** Dec 02, 2024**AT** PADAGIS US **0.05%** **A090898 001** Jun 16, 2011**AT** TARO **0.05%** **A208842 001** Mar 26, 2018**AT** ZYDUS PHARMS **0.05%** **A206378 001** Feb 16, 2017CLOBEX**AT** +! GALDERMA LABS LP **0.05%** **N021835 001** Oct 27, 2005

SUSPENSION/DROPS; OPHTHALMIC

CLOBETASOL PROPIONATE

+! EYENOVIA 0.05% N218158 001 Mar 04, 2024

PRESCRIPTION DRUG PRODUCT LIST

CLOCORTOLONE PIVALATE

CREAM; TOPICAL

CLOCORTOLONE PIVALATE

AB	TARO	<u>0.1%</u>	<u>A206370</u>	<u>001</u>	Apr 21, 2020
-----------	------	-------------	----------------	------------	--------------

CLODERM

AB	+!	LEGACY PHARMA	<u>0.1%</u>	<u>N017765</u>	<u>001</u>
-----------	----	---------------	-------------	----------------	------------

CLOFARABINE

SOLUTION; INTRAVENOUS

CLOFARABINE

AP	ACCORD HLTHCARE	<u>20MG/20ML (1MG/ML)</u>	<u>A212034</u>	<u>001</u>	Feb 22, 2019
-----------	-----------------	---------------------------	----------------	------------	--------------

AP	AMNEAL	<u>20MG/20ML (1MG/ML)</u>	<u>A208857</u>	<u>001</u>	Nov 06, 2017
-----------	--------	---------------------------	----------------	------------	--------------

AP	DR REDDYS	<u>20MG/20ML (1MG/ML)</u>	<u>A205375</u>	<u>001</u>	Nov 06, 2017
-----------	-----------	---------------------------	----------------	------------	--------------

AP	GLAND PHARMA LTD	<u>20MG/20ML (1MG/ML)</u>	<u>A207831</u>	<u>001</u>	Oct 31, 2018
-----------	------------------	---------------------------	----------------	------------	--------------

AP	MEITHEAL	<u>20MG/20ML (1MG/ML)</u>	<u>A213461</u>	<u>001</u>	Oct 23, 2020
-----------	----------	---------------------------	----------------	------------	--------------

AP	MSN	<u>20MG/20ML (1MG/ML)</u>	<u>A209775</u>	<u>001</u>	Dec 06, 2017
-----------	-----	---------------------------	----------------	------------	--------------

AP	MYLAN LABS LTD	<u>20MG/20ML (1MG/ML)</u>	<u>A208860</u>	<u>001</u>	Nov 06, 2017
-----------	----------------	---------------------------	----------------	------------	--------------

AP	SCINOPHARM TAIWAN	<u>20MG/20ML (1MG/ML)</u>	<u>A216233</u>	<u>001</u>	Jan 03, 2025
-----------	-------------------	---------------------------	----------------	------------	--------------

CLOLAR

AP	+!	GENZYME	<u>20MG/20ML (1MG/ML)</u>	<u>N021673</u>	<u>001</u>	Dec 28, 2004
-----------	----	---------	---------------------------	----------------	------------	--------------

CLOMIPHENE CITRATE

TABLET; ORAL

CLOMIPHENE CITRATE

AB	APPCO	<u>50MG</u>	<u>A216739</u>	<u>001</u>	Nov 08, 2024
-----------	-------	-------------	----------------	------------	--------------

AB	!	COSETTE	<u>50MG</u>	<u>A075528</u>	<u>001</u>	Aug 30, 1999
-----------	---	---------	-------------	----------------	------------	--------------

AB		RUBICON	<u>50MG</u>	<u>A216545</u>	<u>001</u>	Nov 12, 2024
-----------	--	---------	-------------	----------------	------------	--------------

CLOMIPRAMINE HYDROCHLORIDE

CAPSULE; ORAL

ANAFRANIL

AB	+!	SPECGX LLC	<u>25MG</u>	<u>N019906</u>	<u>001</u>	Dec 29, 1989
-----------	----	------------	-------------	----------------	------------	--------------

AB	+		<u>50MG</u>	<u>N019906</u>	<u>002</u>	Dec 29, 1989
-----------	---	--	-------------	----------------	------------	--------------

AB	+		<u>75MG</u>	<u>N019906</u>	<u>003</u>	Dec 29, 1989
-----------	---	--	-------------	----------------	------------	--------------

CLOMIPRAMINE HYDROCHLORIDE

AB		ALEMBIC	<u>25MG</u>	<u>A211822</u>	<u>001</u>	Aug 04, 2021
-----------	--	---------	-------------	----------------	------------	--------------

AB			<u>50MG</u>	<u>A211822</u>	<u>002</u>	Aug 04, 2021
-----------	--	--	-------------	----------------	------------	--------------

AB			<u>75MG</u>	<u>A211822</u>	<u>003</u>	Aug 04, 2021
-----------	--	--	-------------	----------------	------------	--------------

AB		AUROBINDO PHARMA	<u>25MG</u>	<u>A216440</u>	<u>001</u>	Mar 20, 2023
-----------	--	------------------	-------------	----------------	------------	--------------

AB			<u>50MG</u>	<u>A216440</u>	<u>002</u>	Mar 20, 2023
-----------	--	--	-------------	----------------	------------	--------------

AB			<u>75MG</u>	<u>A216440</u>	<u>003</u>	Mar 20, 2023
-----------	--	--	-------------	----------------	------------	--------------

AB		CHARTWELL RX	<u>25MG</u>	<u>A074364</u>	<u>001</u>	Mar 29, 1996
-----------	--	--------------	-------------	----------------	------------	--------------

AB			<u>50MG</u>	<u>A074364</u>	<u>002</u>	Mar 29, 1996
-----------	--	--	-------------	----------------	------------	--------------

AB			<u>75MG</u>	<u>A074364</u>	<u>003</u>	Mar 29, 1996
-----------	--	--	-------------	----------------	------------	--------------

AB		CHEMISTRY HLTH	<u>25MG</u>	<u>A211364</u>	<u>001</u>	Feb 07, 2020
-----------	--	----------------	-------------	----------------	------------	--------------

AB			<u>50MG</u>	<u>A211364</u>	<u>002</u>	Feb 07, 2020
-----------	--	--	-------------	----------------	------------	--------------

AB			<u>75MG</u>	<u>A211364</u>	<u>003</u>	Feb 07, 2020
-----------	--	--	-------------	----------------	------------	--------------

AB		JUBILANT CADISTA	<u>25MG</u>	<u>A212218</u>	<u>001</u>	Oct 21, 2019
-----------	--	------------------	-------------	----------------	------------	--------------

AB			<u>50MG</u>	<u>A212218</u>	<u>002</u>	Oct 21, 2019
-----------	--	--	-------------	----------------	------------	--------------

AB			<u>75MG</u>	<u>A212218</u>	<u>003</u>	Oct 21, 2019
-----------	--	--	-------------	----------------	------------	--------------

AB		LUPIN LTD	<u>25MG</u>	<u>A209294</u>	<u>001</u>	Nov 21, 2018
-----------	--	-----------	-------------	----------------	------------	--------------

AB			<u>50MG</u>	<u>A209294</u>	<u>002</u>	Nov 21, 2018
-----------	--	--	-------------	----------------	------------	--------------

AB			<u>75MG</u>	<u>A209294</u>	<u>003</u>	Nov 21, 2018
-----------	--	--	-------------	----------------	------------	--------------

AB		MANKIND PHARMA	<u>25MG</u>	<u>A211767</u>	<u>001</u>	Apr 08, 2019
-----------	--	----------------	-------------	----------------	------------	--------------

AB			<u>50MG</u>	<u>A211767</u>	<u>002</u>	Apr 08, 2019
-----------	--	--	-------------	----------------	------------	--------------

AB			<u>75MG</u>	<u>A211767</u>	<u>003</u>	Apr 08, 2019
-----------	--	--	-------------	----------------	------------	--------------

AB		MICRO LABS	<u>25MG</u>	<u>A213219</u>	<u>001</u>	Jun 22, 2020
-----------	--	------------	-------------	----------------	------------	--------------

AB			<u>50MG</u>	<u>A213219</u>	<u>002</u>	Jun 22, 2020
-----------	--	--	-------------	----------------	------------	--------------

AB			<u>75MG</u>	<u>A213219</u>	<u>003</u>	Jun 22, 2020
-----------	--	--	-------------	----------------	------------	--------------

AB		SANDOZ	<u>25MG</u>	<u>A074953</u>	<u>001</u>	Jun 25, 1997
-----------	--	--------	-------------	----------------	------------	--------------

AB			<u>50MG</u>	<u>A074953</u>	<u>002</u>	Jun 25, 1997
-----------	--	--	-------------	----------------	------------	--------------

AB			<u>75MG</u>	<u>A074953</u>	<u>003</u>	Jun 25, 1997
-----------	--	--	-------------	----------------	------------	--------------

AB		TARO	<u>25MG</u>	<u>A074694</u>	<u>001</u>	Dec 31, 1996
-----------	--	------	-------------	----------------	------------	--------------

AB			<u>50MG</u>	<u>A074694</u>	<u>002</u>	Dec 31, 1996
-----------	--	--	-------------	----------------	------------	--------------

AB			<u>75MG</u>	<u>A074694</u>	<u>003</u>	Dec 31, 1996
-----------	--	--	-------------	----------------	------------	--------------

AB		TRUPHARMA	<u>25MG</u>	<u>A210653</u>	<u>001</u>	Apr 03, 2020
-----------	--	-----------	-------------	----------------	------------	--------------

AB			<u>50MG</u>	<u>A210653</u>	<u>002</u>	Apr 03, 2020
-----------	--	--	-------------	----------------	------------	--------------

AB			<u>75MG</u>	<u>A210653</u>	<u>003</u>	Apr 03, 2020
-----------	--	--	-------------	----------------	------------	--------------

AB		UNIQUE	<u>25MG</u>	<u>A212285</u>	<u>001</u>	Aug 07, 2020
-----------	--	--------	-------------	----------------	------------	--------------

AB			<u>50MG</u>	<u>A212285</u>	<u>002</u>	Aug 07, 2020
-----------	--	--	-------------	----------------	------------	--------------

AB			<u>75MG</u>	<u>A212285</u>	<u>003</u>	Aug 07, 2020
-----------	--	--	-------------	----------------	------------	--------------

AB		ZYDUS PHARMS	<u>25MG</u>	<u>A208961</u>	<u>001</u>	Dec 27, 2017
-----------	--	--------------	-------------	----------------	------------	--------------

AB			<u>50MG</u>	<u>A208961</u>	<u>002</u>	Dec 27, 2017
-----------	--	--	-------------	----------------	------------	--------------

AB			<u>75MG</u>	<u>A208961</u>	<u>003</u>	Dec 27, 2017
-----------	--	--	-------------	----------------	------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

CLONAZEPAM

TABLET; ORAL

CLONAZEPAM

<u>AB</u>	ACCORD HLTHCARE	<u>0.5MG</u>	<u>A077147 001</u>	May 02, 2005
<u>AB</u>		<u>1MG</u>	<u>A077147 002</u>	May 02, 2005
<u>AB</u>		<u>2MG</u>	<u>A077147 003</u>	May 02, 2005
<u>AB</u>	ACTAVIS ELIZABETH	<u>0.5MG</u>	<u>A074869 001</u>	Oct 31, 1996
<u>AB</u>		<u>1MG</u>	<u>A074869 002</u>	Oct 31, 1996
<u>AB</u>		<u>2MG</u>	<u>A074869 003</u>	Oct 31, 1996
<u>AB</u>	AUROBINDO PHARMA USA	<u>0.5MG</u>	<u>A075150 001</u>	Oct 05, 1998
<u>AB</u>		<u>1MG</u>	<u>A075150 002</u>	Oct 05, 1998
<u>AB</u>		<u>2MG</u>	<u>A075150 003</u>	Oct 05, 1998
<u>AB</u>	PRINSTON INC	<u>0.5MG</u>	<u>A077856 001</u>	Jun 28, 2006
<u>AB</u>		<u>1MG</u>	<u>A077856 002</u>	Jun 28, 2006
<u>AB</u>		<u>2MG</u>	<u>A077856 003</u>	Jun 28, 2006
<u>AB</u>	RUBICON	<u>0.5MG</u>	<u>A075468 001</u>	Oct 06, 2000
<u>AB</u>		<u>1MG</u>	<u>A075468 002</u>	Oct 06, 2000
<u>AB</u>		<u>2MG</u>	<u>A075468 003</u>	Oct 06, 2000
<u>AB</u>	SANDOZ	<u>0.5MG</u>	<u>A074979 001</u>	Aug 29, 1997
<u>AB</u>		<u>1MG</u>	<u>A074979 002</u>	Aug 29, 1997
<u>AB</u>		<u>2MG</u>	<u>A074979 003</u>	Aug 29, 1997
<u>AB</u>	TEVA	<u>0.5MG</u>	<u>A074569 001</u>	Sep 10, 1996
<u>AB</u>		<u>1MG</u>	<u>A074569 002</u>	Sep 10, 1996
<u>AB</u>		<u>2MG</u>	<u>A074569 003</u>	Sep 10, 1996

KLONOPIN

<u>AB</u>	+ CHEPLAPHARM	<u>0.5MG</u>	<u>N017533 001</u>	
<u>AB</u>	+!	<u>1MG</u>	<u>N017533 002</u>	
<u>AB</u>	+	<u>2MG</u>	<u>N017533 003</u>	

TABLET, ORALLY DISINTEGRATING; ORAL

CLONAZEPAM

<u>AB</u>	ALEMBIC	<u>0.125MG</u>	<u>A211033 001</u>	Jun 28, 2019
<u>AB</u>		<u>0.25MG</u>	<u>A211033 002</u>	Jun 28, 2019
<u>AB</u>		<u>0.5MG</u>	<u>A211033 003</u>	Jun 28, 2019
<u>AB</u>		<u>1MG</u>	<u>A211033 004</u>	Jun 28, 2019
<u>AB</u>		<u>2MG</u>	<u>A211033 005</u>	Jun 28, 2019
<u>AB</u>	BARR	<u>0.125MG</u>	<u>A077194 001</u>	Aug 10, 2005
<u>AB</u>		<u>0.25MG</u>	<u>A077194 002</u>	Aug 10, 2005
<u>AB</u>		<u>0.5MG</u>	<u>A077194 003</u>	Aug 10, 2005
<u>AB</u>		<u>1MG</u>	<u>A077194 004</u>	Aug 10, 2005
<u>AB</u>		<u>2MG</u>	<u>A077194 005</u>	Aug 10, 2005
<u>AB</u>	ENDO OPERATIONS	<u>0.125MG</u>	<u>A077171 001</u>	Aug 03, 2005
<u>AB</u>		<u>0.25MG</u>	<u>A077171 002</u>	Aug 03, 2005
<u>AB</u>		<u>0.5MG</u>	<u>A077171 003</u>	Aug 03, 2005
<u>AB</u>	!	<u>1MG</u>	<u>A077171 004</u>	Aug 03, 2005
<u>AB</u>		<u>2MG</u>	<u>A077171 005</u>	Aug 03, 2005
<u>AB</u>	SUN PHARM INDS INC	<u>0.125MG</u>	<u>A078654 001</u>	Aug 27, 2014
<u>AB</u>		<u>0.25MG</u>	<u>A078654 002</u>	Aug 27, 2014
<u>AB</u>		<u>0.5MG</u>	<u>A078654 003</u>	Aug 27, 2014
<u>AB</u>		<u>1MG</u>	<u>A078654 004</u>	Aug 27, 2014
<u>AB</u>		<u>2MG</u>	<u>A078654 005</u>	Aug 27, 2014

CLONIDINE

SYSTEM; TRANSDERMAL

CATAPRES-TTS-1

<u>AB</u>	+ LAVIPHARM	<u>0.1MG/24HR</u>	<u>N018891 001</u>	Oct 10, 1984
<u>AB</u>				
<u>AB</u>	+ LAVIPHARM	<u>0.2MG/24HR</u>	<u>N018891 002</u>	Oct 10, 1984
<u>AB</u>				
<u>AB</u>	+! LAVIPHARM	<u>0.3MG/24HR</u>	<u>N018891 003</u>	Oct 10, 1984

CLONIDINE

<u>AB</u>	ACTAVIS LABS UT INC	<u>0.1MG/24HR</u>	<u>A090873 001</u>	May 06, 2014
<u>AB</u>		<u>0.2MG/24HR</u>	<u>A090873 002</u>	May 06, 2014
<u>AB</u>		<u>0.3MG/24HR</u>	<u>A090873 003</u>	May 06, 2014
<u>AB</u>	DIFGEN PHARMS	<u>0.1MG/24HR</u>	<u>A076157 001</u>	Aug 18, 2009
<u>AB</u>		<u>0.2MG/24HR</u>	<u>A076157 002</u>	Aug 18, 2009
<u>AB</u>		<u>0.3MG/24HR</u>	<u>A076157 003</u>	Aug 18, 2009
<u>AB</u>	DR REDDYS LABS SA	<u>0.1MG/24HR</u>	<u>A079090 001</u>	Aug 20, 2010
<u>AB</u>		<u>0.2MG/24HR</u>	<u>A079090 002</u>	Aug 20, 2010
<u>AB</u>		<u>0.3MG/24HR</u>	<u>A079090 003</u>	Aug 20, 2010
<u>AB</u>	MYLAN TECHNOLOGIES	<u>0.1MG/24HR</u>	<u>A076166 001</u>	Jul 16, 2010
<u>AB</u>		<u>0.2MG/24HR</u>	<u>A076166 002</u>	Jul 16, 2010
<u>AB</u>		<u>0.3MG/24HR</u>	<u>A076166 003</u>	Jul 16, 2010

PRESCRIPTION DRUG PRODUCT LIST

CLONIDINE

TABLET, EXTENDED RELEASE;ORAL

NEXICLON XR

ATHENA

EQ 0.17MG BASE

N022500 001 Dec 03, 2009

CLONIDINE HYDROCHLORIDE

INJECTABLE; INJECTION

CLONIDINE HYDROCHLORIDE

<u>AP</u>	FRESENIUS KABI USA	<u>1MG/10ML (0.1MG/ML)</u>	<u>A200673 001</u>	Jul 08, 2011
<u>AP</u>		<u>5MG/10ML (0.5MG/ML)</u>	<u>A200673 002</u>	Jul 08, 2011
<u>AP</u>	HIKMA FARMACEUTICA	<u>1MG/10ML (0.1MG/ML)</u>	<u>A200300 001</u>	Jan 26, 2011
<u>AP</u>	!	<u>5MG/10ML (0.5MG/ML)</u>	<u>A200300 002</u>	Jan 26, 2011
<u>AP</u>	XGEN PHARMS	<u>1MG/10ML (0.1MG/ML)</u>	<u>A203167 001</u>	Oct 29, 2013
<u>AP</u>		<u>5MG/10ML (0.5MG/ML)</u>	<u>A203167 002</u>	Oct 29, 2013
<u>AP</u>	ZYDUS PHARMS	<u>1MG/10ML (0.1MG/ML)</u>	<u>A202601 001</u>	Feb 20, 2014
<u>AP</u>		<u>5MG/10ML (0.5MG/ML)</u>	<u>A202601 002</u>	Feb 20, 2014

DURACLON

<u>AP</u>	+	MYLAN INSTITUTIONAL	<u>1MG/10ML (0.1MG/ML)</u>	<u>N020615 001</u>	Oct 02, 1996
SUSPENSION, EXTENDED RELEASE;ORAL					
ONYDA XR					
	+	!	TRIS PHARMA INC	0.1MG/ML	N217645 001 May 24, 2024
TABLET;ORAL					

CLONIDINE HYDROCHLORIDE

<u>AB</u>	ACTAVIS ELIZABETH	<u>0.1MG</u>	<u>A070974 001</u>	Dec 16, 1986
<u>AB</u>		<u>0.2MG</u>	<u>A070975 001</u>	Dec 16, 1986
<u>AB</u>		<u>0.3MG</u>	<u>A070976 001</u>	Dec 16, 1986
<u>AB</u>	ALEMBIC PHARMS LTD	<u>0.1MG</u>	<u>A091368 001</u>	Dec 06, 2011
<u>AB</u>		<u>0.2MG</u>	<u>A091368 002</u>	Dec 06, 2011
<u>AB</u>		<u>0.3MG</u>	<u>A091368 003</u>	Dec 06, 2011
<u>AB</u>	AUROBINDO PHARMA LTD	<u>0.1MG</u>	<u>A070886 002</u>	Aug 31, 1988
<u>AB</u>		<u>0.2MG</u>	<u>A070886 001</u>	Aug 31, 1988
<u>AB</u>		<u>0.3MG</u>	<u>A070886 003</u>	Aug 31, 1988
<u>AB</u>	CHARTWELL MOLECULES	<u>0.1MG</u>	<u>A071785 002</u>	Apr 05, 1988
<u>AB</u>		<u>0.2MG</u>	<u>A071785 003</u>	Apr 05, 1988
<u>AB</u>		<u>0.3MG</u>	<u>A071785 001</u>	Apr 05, 1988
<u>AB</u>	IMPAX LABS	<u>0.1MG</u>	<u>A078099 001</u>	Aug 27, 2009
<u>AB</u>		<u>0.2MG</u>	<u>A078099 002</u>	Aug 27, 2009
<u>AB</u>		<u>0.3MG</u>	<u>A078099 003</u>	Aug 27, 2009
<u>AB</u>	PRINSTON INC	<u>0.1MG</u>	<u>A077901 001</u>	Mar 09, 2007
<u>AB</u>		<u>0.2MG</u>	<u>A077901 002</u>	Mar 09, 2007
<u>AB</u>		<u>0.3MG</u>	<u>A077901 003</u>	Mar 09, 2007
<u>AB</u>	TRUPHARMA	<u>0.1MG</u>	<u>A070923 003</u>	Sep 04, 1987
<u>AB</u>		<u>0.2MG</u>	<u>A070923 002</u>	Sep 04, 1987
<u>AB</u>		<u>0.3MG</u>	<u>A070923 001</u>	Sep 04, 1987
<u>AB</u>	UNICHEM	<u>0.1MG</u>	<u>A078895 001</u>	Aug 26, 2009
<u>AB</u>		<u>0.2MG</u>	<u>A078895 002</u>	Aug 26, 2009
<u>AB</u>	!	<u>0.3MG</u>	<u>A078895 003</u>	Aug 26, 2009
<u>AB</u>	YUNG SHIN PHARM	<u>0.1MG</u>	<u>A202297 001</u>	Jun 13, 2013
<u>AB</u>		<u>0.2MG</u>	<u>A202297 002</u>	Jun 13, 2013
<u>AB</u>		<u>0.3MG</u>	<u>A202297 003</u>	Jun 13, 2013

TABLET, EXTENDED RELEASE;ORAL

CLONIDINE HYDROCHLORIDE

<u>AB1</u>	ACTAVIS ELIZABETH	<u>0.1MG</u>	<u>A203320 001</u>	May 15, 2015	
<u>AB1</u>	!	AJANTA PHARMA LTD	<u>0.1MG</u>	<u>A209686 001</u>	Nov 20, 2017
<u>AB1</u>	CHARTWELL RX	<u>0.1MG</u>	<u>A209285 001</u>	Oct 23, 2017	
<u>AB1</u>	JUBILANT GENERICS	<u>0.1MG</u>	<u>A210338 001</u>	Jan 29, 2018	
<u>AB1</u>	NOVAST LABS	<u>0.1MG</u>	<u>A209675 001</u>	Mar 05, 2019	
<u>AB1</u>	XIAMEN LP PHARM CO	<u>0.1MG</u>	<u>A209757 001</u>	Nov 20, 2017	

CLOPIDOGREL BISULFATE

TABLET;ORAL

CLOPIDOGREL BISULFATE

<u>AB</u>	ACME LABS	<u>EQ 75MG BASE</u>	<u>A078004 001</u>	May 17, 2012
<u>AB</u>	ALKEM LABS LTD	<u>EQ 75MG BASE</u>	<u>A203632 001</u>	Dec 08, 2023
<u>AB</u>	AMNEAL PHARMS	<u>EQ 75MG BASE</u>	<u>A203751 001</u>	Apr 11, 2014
<u>AB</u>	APOTEX INC	<u>EQ 75MG BASE</u>	<u>A076274 001</u>	May 17, 2012
<u>AB</u>	!	<u>EQ 300MG BASE</u>	<u>A076274 002</u>	Mar 04, 2014
<u>AB</u>	AUROBINDO PHARMA LTD	<u>EQ 75MG BASE</u>	<u>A090540 001</u>	May 17, 2012
<u>AB</u>	DR REDDYS	<u>EQ 75MG BASE</u>	<u>A076273 001</u>	Jan 14, 2008
<u>AB</u>	DR REDDYS LABS LTD	<u>EQ 75MG BASE</u>	<u>A091023 002</u>	Aug 05, 2024
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A091023 001</u>	May 17, 2012
<u>AB</u>	HETERO LABS LTD V	<u>EQ 75MG BASE</u>	<u>A205345 001</u>	Aug 04, 2023

PRESCRIPTION DRUG PRODUCT LIST

CLOPIDOGREL BISULFATE

TABLET; ORAL

CLOPIDOGREL BISULFATE

<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A205345 002</u>	Aug 04, 2023
<u>AB</u>	MACLEODS PHARMS LTD	<u>EQ 75MG BASE</u>	<u>A202928 001</u>	Feb 10, 2014
<u>AB</u>	MSN	<u>EQ 75MG BASE</u>	<u>A215388 001</u>	Sep 20, 2024
<u>AB</u>	POLYGEN PHARMS	<u>EQ 75MG BASE</u>	<u>A213351 001</u>	Jul 17, 2020
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A213351 002</u>	Jul 17, 2020
<u>AB</u>	PRINSTON INC	<u>EQ 75MG BASE</u>	<u>A206376 001</u>	May 07, 2018
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A206376 002</u>	May 07, 2018
<u>AB</u>	RISING	<u>EQ 75MG BASE</u>	<u>A204359 001</u>	Feb 02, 2017
<u>AB</u>	SCIEGEN PHARMS INC	<u>EQ 75MG BASE</u>	<u>A204165 001</u>	Sep 15, 2014
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A204165 002</u>	Sep 15, 2014
<u>AB</u>	TEVA	<u>EQ 75MG BASE</u>	<u>A076999 001</u>	May 17, 2012
<u>AB</u>	TORRENT PHARMS LTD	<u>EQ 75MG BASE</u>	<u>A090844 001</u>	May 17, 2012

PLAVIX

<u>AB</u>	+	SANOFI AVENTIS US	<u>EQ 75MG BASE</u>	<u>N020839 001</u>	Nov 17, 1997
<u>AB</u>	+		<u>EQ 300MG BASE</u>	<u>N020839 002</u>	Sep 20, 2007

CLORAZEPATE DIPOTASSIUM

TABLET; ORAL

CLORAZEPATE DIPOTASSIUM

<u>AB</u>	AUROBINDO PHARMA	<u>3.75MG</u>	<u>A071858 002</u>	Jul 17, 1987
<u>AB</u>		<u>7.5MG</u>	<u>A071858 003</u>	Jul 17, 1987
<u>AB</u>	!	<u>15MG</u>	<u>A071858 001</u>	Jul 17, 1987
<u>AB</u>	COREPHARMA	<u>3.75MG</u>	<u>A215566 001</u>	Jun 14, 2022
<u>AB</u>		<u>7.5MG</u>	<u>A215566 002</u>	Jun 14, 2022
<u>AB</u>		<u>15MG</u>	<u>A215566 003</u>	Jun 14, 2022
<u>AB</u>	NOVITIUM PHARMA	<u>3.75MG</u>	<u>A213730 001</u>	Jun 16, 2022
<u>AB</u>		<u>7.5MG</u>	<u>A213730 002</u>	Jun 16, 2022
<u>AB</u>		<u>15MG</u>	<u>A213730 003</u>	Dec 16, 2022
<u>AB</u>	TARO	<u>3.75MG</u>	<u>A075731 003</u>	Apr 27, 2000
<u>AB</u>		<u>7.5MG</u>	<u>A075731 002</u>	Apr 27, 2000
<u>AB</u>		<u>15MG</u>	<u>A075731 001</u>	Apr 27, 2000

TRANKENE

<u>AB</u>	+	AJENAT PHARMS	<u>7.5MG</u>	<u>N017105 007</u>	
-----------	---	---------------	--------------	--------------------	--

CLOTTRIMAZOLE

CREAM; TOPICAL

CLOTTRIMAZOLE

<u>AB</u>	FOUGERA PHARMS	<u>1%</u>	<u>A078338 001</u>	Sep 02, 2008	
<u>AB</u>	GLENMARK PHARMS	<u>1%</u>	<u>A090219 001</u>	Aug 03, 2010	
<u>AB</u>	!	TARO	<u>1%</u>	<u>A072640 001</u>	Aug 31, 1993

SOLUTION; TOPICAL

CLOTTRIMAZOLE

<u>AT</u>	NOVITIUM PHARMA	<u>1%</u>	<u>A209815 001</u>	Feb 14, 2019	
<u>AT</u>	SCIEGEN PHARMS INC	<u>1%</u>	<u>A216569 001</u>	Oct 16, 2023	
<u>AT</u>	!	TARO	<u>1%</u>	<u>A074580 001</u>	Jul 29, 1996
<u>AT</u>	TEVA	<u>1%</u>	<u>A073306 001</u>	Feb 28, 1995	
<u>AT</u>	TRUPHARMA	<u>1%</u>	<u>A212281 001</u>	Jul 25, 2019	

TROCHE/LOZENGE; ORAL

CLOTTRIMAZOLE

<u>AB</u>	!	HIKMA	<u>10MG</u>	<u>A076387 001</u>	Jul 29, 2004
<u>AB</u>		PADAGIS US	<u>10MG</u>	<u>A076763 001</u>	Oct 28, 2005
<u>AB</u>		THINQ PHARM-CRO PVT	<u>10MG</u>	<u>A215641 001</u>	Feb 29, 2024

CLOZAPINE

SUSPENSION; ORAL

VERSACLOZ

+	!	DOUGLAS PHARMS	50MG/ML	N203479 001	Feb 06, 2013
---	---	----------------	---------	-------------	--------------

TABLET; ORAL

CLOZAPINE

<u>AB</u>	ACCORD HLTHCARE	<u>25MG</u>	<u>A202873 001</u>	Nov 25, 2015
<u>AB</u>		<u>100MG</u>	<u>A202873 002</u>	Nov 25, 2015
<u>AB</u>	AUROBINDO PHARMA	<u>25MG</u>	<u>A206433 001</u>	Nov 29, 2016
<u>AB</u>		<u>50MG</u>	<u>A206433 002</u>	Nov 29, 2016
<u>AB</u>		<u>100MG</u>	<u>A206433 003</u>	Nov 29, 2016
<u>AB</u>		<u>200MG</u>	<u>A206433 004</u>	Nov 29, 2016
<u>AB</u>	IVAX SUB TEVA PHARMS	<u>25MG</u>	<u>A074949 001</u>	Nov 26, 1997
<u>AB</u>		<u>50MG</u>	<u>A074949 004</u>	Apr 25, 2005
<u>AB</u>		<u>50MG</u>	<u>A076809 003</u>	Dec 16, 2005
<u>AB</u>		<u>100MG</u>	<u>A074949 002</u>	Nov 26, 1997
<u>AB</u>		<u>100MG</u>	<u>A076809 002</u>	Dec 16, 2005

PRESCRIPTION DRUG PRODUCT LIST

CLOZAPINE

TABLET; ORAL

CLOZAPINE

<u>AB</u>		<u>200MG</u>	<u>A076809 001</u>	Dec 16, 2005
<u>AB</u>	MYLAN	<u>25MG</u>	<u>A075417 001</u>	May 27, 1999
<u>AB</u>		<u>50MG</u>	<u>A075417 004</u>	Apr 15, 2010
<u>AB</u>		<u>100MG</u>	<u>A075417 002</u>	May 27, 1999
<u>AB</u>		<u>200MG</u>	<u>A075417 005</u>	Apr 15, 2010
<u>AB</u>	SUN PHARM INDS INC	<u>25MG</u>	<u>A075713 001</u>	Nov 15, 2002
<u>AB</u>		<u>50MG</u>	<u>A075713 003</u>	Aug 19, 2005
<u>AB</u>		<u>100MG</u>	<u>A075713 002</u>	Nov 15, 2002
<u>AB</u>		<u>200MG</u>	<u>A075713 004</u>	Nov 07, 2017

CLOZARIL

<u>AB</u>	+	HERITAGE LIFE	<u>25MG</u>	<u>N019758 001</u>	Sep 26, 1989
<u>AB</u>	+	!	<u>100MG</u>	<u>N019758 002</u>	Sep 26, 1989

CLOZAPINE

	IVAX SUB TEVA	12.5MG	A074949 003	Jul 31, 2003
--	---------------	--------	-------------	--------------

PHARMS

TABLET, ORALLY DISINTEGRATING; ORAL

CLOZAPINE

<u>AB</u>	AUROBINDO PHARMA	<u>12.5MG</u>	<u>A212923 001</u>	Dec 12, 2024
<u>AB</u>		<u>25MG</u>	<u>A212923 002</u>	Dec 12, 2024
<u>AB</u>		<u>100MG</u>	<u>A212923 003</u>	Dec 12, 2024
<u>AB</u>		<u>150MG</u>	<u>A212923 004</u>	Dec 12, 2024
<u>AB</u>		<u>200MG</u>	<u>A212923 005</u>	Dec 12, 2024
<u>AB</u>	BARR LABS INC	<u>12.5MG</u>	<u>A090308 003</u>	Apr 09, 2018
<u>AB</u>		<u>25MG</u>	<u>A090308 001</u>	Nov 25, 2015
<u>AB</u>	!	<u>100MG</u>	<u>A090308 002</u>	Nov 25, 2015
<u>AB</u>		<u>150MG</u>	<u>A090308 004</u>	Nov 25, 2015
<u>AB</u>		<u>200MG</u>	<u>A090308 005</u>	Nov 25, 2015
<u>AB</u>	MYLAN	<u>25MG</u>	<u>A201824 002</u>	Sep 15, 2015
<u>AB</u>		<u>100MG</u>	<u>A201824 003</u>	Sep 15, 2015
<u>AB</u>		<u>150MG</u>	<u>A201824 004</u>	Aug 07, 2023
<u>AB</u>		<u>200MG</u>	<u>A201824 005</u>	Aug 07, 2023

COBICISTAT

TABLET; ORAL

TYBOST

+	!	GILEAD SCIENCES INC	150MG	N203094 001	Sep 24, 2014
---	---	---------------------	-------	-------------	--------------

COBICISTAT; DARUNAVIR

TABLET; ORAL

PREZCOBIX

+	!	JANSSEN PRODS	150MG; 800MG	N205395 001	Jan 29, 2015
---	---	---------------	--------------	-------------	--------------

COBICISTAT; DARUNAVIR; EMTRICITABINE; TENOFOVIR ALAFENAMIDE FUMARATE

TABLET; ORAL

SYM TUZA

+	!	JANSSEN PRODS	150MG; 800MG; 200MG; EQ 10MG BASE	N210455 001	Jul 17, 2018
---	---	---------------	-----------------------------------	-------------	--------------

COBICISTAT; ELVITEGRAVIR; EMTRICITABINE; TENOFOVIR ALAFENAMIDE FUMARATE

TABLET; ORAL

GENVOYA

+	!	GILEAD SCIENCES INC	150MG; 150MG; 200MG; EQ 10MG BASE	N207561 001	Nov 05, 2015
---	---	---------------------	-----------------------------------	-------------	--------------

COBICISTAT; ELVITEGRAVIR; EMTRICITABINE; TENOFOVIR DISOPROXIL FUMARATE

TABLET; ORAL

STRIBILD

+	!	GILEAD SCIENCES INC	150MG; 150MG; 200MG; 300MG	N203100 001	Aug 27, 2012
---	---	---------------------	----------------------------	-------------	--------------

COBIMETINIB FUMARATE

TABLET; ORAL

COTELLIC

+	!	GENENTECH INC	EQ 20MG BASE	N206192 001	Nov 10, 2015
---	---	---------------	--------------	-------------	--------------

COCAINE HYDROCHLORIDE

SOLUTION; NASAL

GOPRELTO

+	!	NODEN PHARMA	4%	N209963 001	Dec 14, 2017
---	---	--------------	----	-------------	--------------

NUMBRINO

+	!	OMNIVIUM PHARMS	4%	N209575 001	Jan 10, 2020
---	---	-----------------	----	-------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

CODEINE PHOSPHATE; PHENYLEPHRINE HYDROCHLORIDE; PROMETHAZINE HYDROCHLORIDE

SYRUP;ORAL

PROMETH HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE W/CODEINE PHOSPHATE

AA	!	GENUS	<u>10MG/5ML; 5MG/5ML; 6.25MG/5ML</u>	<u>A040660</u>	<u>001</u>	Dec 07, 2006
AA		AMNEAL PHARMS	<u>10MG/5ML; 5MG/5ML; 6.25MG/5ML</u>	<u>A200963</u>	<u>001</u>	Aug 26, 2015

CODEINE PHOSPHATE; PROMETHAZINE HYDROCHLORIDE

SYRUP;ORAL

PROMETHAZINE HYDROCHLORIDE AND CODEINE PHOSPHATE

AA		AMNEAL PHARMS	<u>10MG/5ML; 6.25MG/5ML</u>	<u>A200894</u>	<u>001</u>	Apr 24, 2013
AA		NOSTRUM LABS INC	<u>10MG/5ML; 6.25MG/5ML</u>	<u>A090180</u>	<u>001</u>	Mar 17, 2010
AA		QUAGEN	<u>10MG/5ML; 6.25MG/5ML</u>	<u>A214238</u>	<u>001</u>	Oct 08, 2020
AA	!	TRIS PHARMA INC	<u>10MG/5ML; 6.25MG/5ML</u>	<u>A200386</u>	<u>001</u>	Jun 29, 2012
AA		WOCKHARDT BIO AG	<u>10MG/5ML; 6.25MG/5ML</u>	<u>A088875</u>	<u>001</u>	Dec 17, 1984

PROMETHAZINE WITH CODEINE

AA		PHARM ASSOC	<u>10MG/5ML; 6.25MG/5ML</u>	<u>A040650</u>	<u>001</u>	Jan 31, 2006
-----------	--	-------------	-----------------------------	----------------	------------	--------------

CODEINE PHOSPHATE; PSEUDOEPHEDRINE HYDROCHLORIDE; TRIPROLIDINE HYDROCHLORIDE

SYRUP;ORAL

TRIACIN-C

!	ANIMA	10MG/5ML; 30MG/5ML; 1.25MG/5ML	A088704	001	Mar 22, 1985
---	-------	--------------------------------	---------	-----	--------------

CODEINE SULFATE

TABLET;ORAL

CODEINE SULFATE

AB	+	HIKMA	<u>15MG</u>	<u>N022402</u>	<u>001</u>	Jul 16, 2009
AB	+		<u>30MG</u>	<u>N022402</u>	<u>002</u>	Jul 16, 2009
AB	+	!	<u>60MG</u>	<u>N022402</u>	<u>003</u>	Jul 16, 2009
AB		LANNETT CO INC	<u>15MG</u>	<u>A203046</u>	<u>001</u>	Jun 13, 2014
AB			<u>30MG</u>	<u>A203046</u>	<u>002</u>	Jun 13, 2014
AB			<u>60MG</u>	<u>A203046</u>	<u>003</u>	Jun 13, 2014

COLCHICINE

CAPSULE;ORAL

COLCHICINE

AB		ANNORA PHARMA	<u>0.6MG</u>	<u>A217620</u>	<u>001</u>	Apr 29, 2024
AB		AUROBINDO PHARMA LTD	<u>0.6MG</u>	<u>A215463</u>	<u>001</u>	Apr 29, 2024
AB		ENDO OPERATIONS	<u>0.6MG</u>	<u>A208678</u>	<u>001</u>	Nov 29, 2018
AB		GRANULES	<u>0.6MG</u>	<u>A210757</u>	<u>001</u>	Apr 29, 2024

MITIGARE

AB	+	!	HIKMA INTL PHARMS	<u>0.6MG</u>	<u>N204820</u>	<u>001</u>	Sep 26, 2014
-----------	---	---	-------------------	--------------	----------------	------------	--------------

SOLUTION;ORAL

GLOPERBA

+	!	SCILEX PHARMS	0.6MG/5ML	N210942	001	Jan 30, 2019
---	---	---------------	-----------	---------	-----	--------------

TABLET;ORAL

COLCHICINE

AB		ALKEM LABS LTD	<u>0.6MG</u>	<u>A211250</u>	<u>001</u>	Feb 08, 2019
AB		AMNEAL PHARMS	<u>0.6MG</u>	<u>A204711</u>	<u>001</u>	Sep 28, 2016
AB		AUROBINDO PHARMA LTD	<u>0.6MG</u>	<u>A215444</u>	<u>001</u>	Jan 06, 2022
AB		DR REDDYS	<u>0.6MG</u>	<u>A209876</u>	<u>001</u>	Sep 06, 2019
AB	!	ENDO OPERATIONS	<u>0.6MG</u>	<u>A203976</u>	<u>001</u>	Aug 12, 2021
AB		GRANULES	<u>0.6MG</u>	<u>A210425</u>	<u>001</u>	Feb 05, 2020
AB		HETERO LABS LTD V	<u>0.6MG</u>	<u>A208993</u>	<u>001</u>	Aug 13, 2021
AB		MYLAN	<u>0.6MG</u>	<u>A209470</u>	<u>001</u>	Sep 16, 2019
AB		STRIDES PHARMA	<u>0.6MG</u>	<u>A209173</u>	<u>001</u>	Mar 10, 2022
AB		WATSON LABS INC	<u>0.6MG</u>	<u>A204461</u>	<u>001</u>	Jul 31, 2019
AB		ZYDUS PHARMS	<u>0.6MG</u>	<u>A211519</u>	<u>001</u>	Feb 19, 2019
			0.3MG	A211519	002	Nov 14, 2019

LODOCO

+	!	AGEPHA PHARMA FZ	0.5MG	N215727	001	Jun 16, 2023
---	---	------------------	-------	---------	-----	--------------

COLCHICINE; PROBENECID

TABLET;ORAL

COL-PROBENECID

AB	+	!	WATSON LABS	<u>0.5MG; 500MG</u>	<u>A084279</u>	<u>001</u>	
AB			NOVAST LABS	<u>0.5MG; 500MG</u>	<u>A040618</u>	<u>001</u>	May 13, 2008
AB			RISING	<u>0.5MG; 500MG</u>	<u>A217030</u>	<u>001</u>	Oct 24, 2023

PRESCRIPTION DRUG PRODUCT LIST

COLESEVELAM HYDROCHLORIDE

FOR SUSPENSION; ORAL

COLESEVELAM HYDROCHLORIDE

AB	ALKEM LABS LTD	3.75GM/PACKET	A210316 001	May 06, 2019
AB	GLENMARK PHARMS LTD	3.75GM/PACKET	A202190 002	Jul 16, 2018

WELCHOL

AB	+ ! COSETTE	3.75GM/PACKET	N022362 002	Oct 02, 2009
	COLESEVELAM HYDROCHLORIDE			
	GLENMARK PHARMS LTD	1.875GM/PACKET	A202190 001	Jul 16, 2018

TABLET; ORAL

COLESEVELAM HYDROCHLORIDE

AB	ALKEM LABS LTD	625MG	A209038 001	Oct 05, 2018
AB	BEIJING TIDE PHARM	625MG	A206036 001	Oct 14, 2021
AB	BIONPHARMA	625MG	A208670 001	Sep 13, 2019
AB	DR REDDYS	625MG	A210889 001	Oct 05, 2018
AB	GLENMARK PHARMS LTD	625MG	A203480 001	May 18, 2018
AB	IMPAX LABS INC	625MG	A091600 001	May 16, 2018
AB	INVENTIA	625MG	A212050 001	Dec 04, 2020
AB	ZHEJIANG JINGXIN	625MG	A209946 001	Jul 15, 2020
AB	ZYDUS PHARMS	625MG	A207765 001	Oct 07, 2019

WELCHOL

AB	+ ! COSETTE	625MG	N021176 001	May 26, 2000
-----------	--------------------	--------------	--------------------	--------------

COLESTIPOL HYDROCHLORIDE

GRANULE; ORAL

COLESTID

AB	+ PFIZER	5GM/SCOOPFUL	N017563 003	Sep 22, 1995
AB	+ !	5GM/PACKET	N017563 004	Sep 22, 1995

COLESTIPOL HYDROCHLORIDE

AB	IMPAX LABS	5GM/SCOOPFUL	A077277 001	May 02, 2006
AB		5GM/PACKET	A077277 002	May 02, 2006

TABLET; ORAL

COLESTID

AB	+ PFIZER	1GM	N020222 001	Jul 19, 1994
-----------	-----------------	------------	--------------------	--------------

COLESTIPOL HYDROCHLORIDE

AB	ANI PHARMS	1GM	A216517 001	Mar 10, 2023
AB	EDENBRIDGE PHARMS	1GM	A217667 001	Aug 16, 2024
AB	! IMPAX LABS	1GM	A077510 001	Oct 24, 2006
AB	RICONPHARMA LLC	1GM	A217462 001	Dec 03, 2024
AB	ZYDUS PHARMS	1GM	A215223 001	Mar 10, 2022

COLISTIMETHATE SODIUM

INJECTABLE; INJECTION

COLISTIMETHATE SODIUM

AP	ADRASTEIA PHARMA	EQ 150MG BASE/VIAL	A064216 001	Feb 26, 1999
AP	AVET LIFESCIENCES	EQ 150MG BASE/VIAL	A202359 001	Sep 28, 2012
AP	FRESENIUS KABI USA	EQ 150MG BASE/VIAL	A065364 001	Apr 17, 2008
AP	NEXUS	EQ 150MG BASE/VIAL	A065177 001	Mar 19, 2004
AP	SAGENT PHARMS INC	EQ 150MG BASE/VIAL	A201365 001	Feb 19, 2014
AP	XELLIA PHARMS APS	EQ 150MG BASE/VIAL	A205356 001	May 29, 2015

COLY-MYCIN M

AP	+ ! ENDO OPERATIONS	EQ 150MG BASE/VIAL	N050108 002	
-----------	----------------------------	---------------------------	--------------------	--

COLISTIN SULFATE; HYDROCORTISONE ACETATE; NEOMYCIN SULFATE; THONZONIUM BROMIDE

SUSPENSION/DROPS; OTIC

COLY-MYCIN S

+ !	ENDO OPERATIONS	EQ 3MG BASE/ML; 10MG/ML; EQ 3.3MG BASE/ML; 0.5MG/ML	N050356 001	
------------	-----------------	--	-------------	--

CONIVAPTAN HYDROCHLORIDE

INJECTABLE; INTRAVENOUS

VAPRISOL IN 5% DEXTROSE IN PLASTIC CONTAINER

+ !	CUMBERLAND	20MG/100ML (0.2MG/ML)	N021697 002	Oct 08, 2008
------------	------------	-----------------------	-------------	--------------

COPPER

SYSTEM; INTRAUTERINE

PARAGARD T 380A

+ !	COOPERSURGICAL	309MG/COPPER	N018680 001	Nov 15, 1984
------------	----------------	--------------	-------------	--------------

COPPER CU-64 DOTATATE

SOLUTION; INTRAVENOUS

DETECTNET

+ !	RADIOMEDIX	4mL (1mCi/ML)	N213227 001	Sep 03, 2020
------------	------------	---------------	-------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

CORTICOTROPIN

INJECTABLE; INJECTION

ACTHAR GEL

+	!	MALLINCKRODT	80 UNITS/ML	N008372	008	
		IRELAND				

ACTHAR GEL (AUTOINJECTOR)

+	!	MALLINCKRODT	40 UNITS/0.5ML	N008372	003	Feb 29, 2024
		IRELAND				

+	!		80 UNITS/ML	N008372	004	Feb 29, 2024
---	---	--	-------------	---------	-----	--------------

PURIFIED CORTROPHIN GEL

+	!	ANI PHARMS	80 UNITS/ML	N008975	002	
---	---	------------	-------------	---------	-----	--

COSYNTROPIN

INJECTABLE; INJECTION

CORTROSYN

<u>AP</u>	+	!	AMPHASTAR PHARMS	<u>0.25MG/VIAL</u>	<u>N016750</u>	<u>001</u>	
			INC				

COSYNTROPIN

<u>AP</u>			SANDOZ	<u>0.25MG/VIAL</u>	<u>A202147</u>	<u>001</u>	Jun 29, 2012
------------------	--	--	--------	---------------------------	-----------------------	-------------------	--------------

CRINECERFONT

CAPSULE; ORAL

CRENESSITY

+		NEUROCRINE	25MG	N218808	001	Dec 13, 2024
---	--	------------	------	---------	-----	--------------

+			50MG	N218808	002	Dec 13, 2024
---	--	--	------	---------	-----	--------------

+	!		100MG	N218808	003	Dec 13, 2024
---	---	--	-------	---------	-----	--------------

SOLUTION; ORAL

CRENESSITY

+	!	NEUROCRINE	50MG/ML	N218820	001	Dec 13, 2024
---	---	------------	---------	---------	-----	--------------

CRISABOROLE

OINTMENT; TOPICAL

EUCRISA

+	!	ANACOR PHARMS INC	2%	N207695	001	Dec 14, 2016
---	---	-------------------	----	---------	-----	--------------

CRIZOTINIB

CAPSULE; ORAL

XALKORI

+		PF PRISM CV	200MG	N202570	001	Aug 26, 2011
---	--	-------------	-------	---------	-----	--------------

+	!		250MG	N202570	002	Aug 26, 2011
---	---	--	-------	---------	-----	--------------

CAPSULE, PELLETS; ORAL

XALKORI

+		PF PRISM CV	20MG	N217581	001	Sep 07, 2023
---	--	-------------	------	---------	-----	--------------

+			50MG	N217581	002	Sep 07, 2023
---	--	--	------	---------	-----	--------------

+	!		150MG	N217581	003	Sep 07, 2023
---	---	--	-------	---------	-----	--------------

CROFELEMER

TABLET, DELAYED RELEASE; ORAL

MYTESI

+	!	NAPO PHARMS INC	125MG	N202292	001	Dec 31, 2012
---	---	-----------------	-------	---------	-----	--------------

CROMOLYN SODIUM

CONCENTRATE; ORAL

CROMOLYN SODIUM

<u>AA</u>			AILEX PHARMS LLC	<u>100MG/5ML</u>	<u>A209264</u>	<u>001</u>	Oct 16, 2017
------------------	--	--	------------------	-------------------------	-----------------------	-------------------	--------------

<u>AA</u>			MICRO LABS LTD	<u>100MG/5ML</u>	<u>A202745</u>	<u>001</u>	Apr 04, 2013
------------------	--	--	----------------	-------------------------	-----------------------	-------------------	--------------

INDIA

<u>AA</u>			RISING	<u>100MG/5ML</u>	<u>A202583</u>	<u>001</u>	Oct 27, 2011
------------------	--	--	--------	-------------------------	-----------------------	-------------------	--------------

GASTROCROM

<u>AA</u>	+	!	MYLAN SPECIALITY LP	<u>100MG/5ML</u>	<u>N020479</u>	<u>001</u>	Feb 29, 1996
------------------	---	---	---------------------	-------------------------	-----------------------	-------------------	--------------

SOLUTION; INHALATION

CROMOLYN SODIUM

<u>AN</u>			AILEX PHARMS LLC	<u>10MG/ML</u>	<u>A209453</u>	<u>001</u>	Oct 16, 2017
------------------	--	--	------------------	-----------------------	-----------------------	-------------------	--------------

<u>AN</u>			MICRO LABS	<u>10MG/ML</u>	<u>A213658</u>	<u>001</u>	Apr 29, 2022
------------------	--	--	------------	-----------------------	-----------------------	-------------------	--------------

<u>AN</u>	!		TEVA PHARMS	<u>10MG/ML</u>	<u>A075271</u>	<u>001</u>	Jan 18, 2000
------------------	---	--	-------------	-----------------------	-----------------------	-------------------	--------------

<u>AN</u>			VIRTUS	<u>10MG/ML</u>	<u>A075437</u>	<u>001</u>	Apr 21, 2000
------------------	--	--	--------	-----------------------	-----------------------	-------------------	--------------

<u>AN</u>			WOCKHARDT BIO AG	<u>10MG/ML</u>	<u>A075346</u>	<u>001</u>	Oct 25, 1999
------------------	--	--	------------------	-----------------------	-----------------------	-------------------	--------------

SOLUTION/DROPS; OPHTHALMIC

CROMOLYN SODIUM

!			SANDOZ	4%	A075282	001	Jun 16, 1999
---	--	--	--------	----	---------	-----	--------------

PRESCRIPTION DRUG PRODUCT LIST

CROTAMITON

LOTION; TOPICAL

CROTAN

AT	LEGACY PHARMA	10%	A087204	001	
-----------	---------------	------------	----------------	------------	--

EURAX

AT	+	JOURNEY	10%	N009112	003
-----------	---	---------	------------	----------------	------------

CUPRIC CHLORIDE

INJECTABLE; INJECTION

CUPRIC CHLORIDE

AP	EXELA PHARMA	EQ 0.4MG COPPER/ML	A212071	001	Oct 31, 2022
-----------	--------------	---------------------------	----------------	------------	--------------

AP	RK PHARMA	EQ 0.4MG COPPER/ML	A217626	001	Apr 30, 2024
-----------	-----------	---------------------------	----------------	------------	--------------

AP	SOMERSET	EQ 0.4MG COPPER/ML	A216113	001	Apr 30, 2024
-----------	----------	---------------------------	----------------	------------	--------------

CUPRIC CHLORIDE IN PLASTIC CONTAINER

AP	+	HOSPIRA	EQ 0.4MG COPPER/ML	N018960	001	Jun 26, 1986
-----------	---	---------	---------------------------	----------------	------------	--------------

CUPRIC SULFATE

INJECTABLE; INJECTION

CUPRIC SULFATE

AP	!	AM REGENT	EQ 0.4MG COPPER/ML	A216324	001	Dec 16, 2022
-----------	---	-----------	---------------------------	----------------	------------	--------------

AP		APOTEX CORP	EQ 0.4MG COPPER/ML	A218745	001	Aug 19, 2024
-----------	--	-------------	---------------------------	----------------	------------	--------------

CUPRIC SULFATE; MANGANESE SULFATE; SELENIOUS ACID; ZINC SULFATE

SOLUTION; INTRAVENOUS

MULTRYS

+	AM REGENT	EQ 60MCG COPPER/ML;EQ 3MCG BASE/ML;EQ 6MCG SELENIUM/ML;EQ 1000MCG BASE/ML (1ML)	N209376	003	Jun 30, 2021
---	-----------	---	---------	-----	--------------

TRALEMENT

+	AM REGENT	EQ 0.3MG COPPER/ML;EQ 55MCG BASE/ML;EQ 60MCG SELENIUM/ML;EQ 3MG BASE/ML (1ML)	N209376	001	Jul 02, 2020
---	-----------	---	---------	-----	--------------

+		EQ 0.3MG COPPER/ML;EQ 55MCG BASE/ML;EQ 60MCG SELENIUM/ML;EQ 3MG BASE/ML (5ML)	N209376	002	Dec 02, 2020
---	--	---	---------	-----	--------------

CYANOCOBALAMIN

INJECTABLE; INJECTION

CYANOCOBALAMIN

AP	+	AM REGENT	1MG/ML	A080737	001	
-----------	---	-----------	---------------	----------------	------------	--

AP		GLAND PHARMA LTD	1MG/ML	A214390	001	Sep 24, 2020
-----------	--	------------------	---------------	----------------	------------	--------------

AP		MANKIND PHARMA	1MG/ML	A217839	001	Jul 10, 2024
-----------	--	----------------	---------------	----------------	------------	--------------

AP		MYLAN LABS LTD	1MG/ML	A204829	001	Jun 05, 2017
-----------	--	----------------	---------------	----------------	------------	--------------

AP		NANJING KING-FRIEND	1MG/ML	A214316	001	Nov 15, 2024
-----------	--	---------------------	---------------	----------------	------------	--------------

AP		SAGENT PHARMS INC	1MG/ML	A215107	001	Nov 15, 2022
-----------	--	-------------------	---------------	----------------	------------	--------------

AP		SANDOZ	1MG/ML	A212915	001	Jan 04, 2021
-----------	--	--------	---------------	----------------	------------	--------------

AP		SOLA PHARMS	1MG/ML	A215417	001	Apr 15, 2022
-----------	--	-------------	---------------	----------------	------------	--------------

AP		SOMERSET THERAPS LLC	1MG/ML	A206503	001	Dec 11, 2015
-----------	--	----------------------	---------------	----------------	------------	--------------

AP			1MG/ML	A209429	001	Dec 18, 2018
-----------	--	--	---------------	----------------	------------	--------------

AP		VITRUVIAS THERAP	1MG/ML	A209255	001	Dec 18, 2018
-----------	--	------------------	---------------	----------------	------------	--------------

AP		WEST-WARD PHARMS	1MG/ML	A080515	002	
-----------	--	------------------	---------------	----------------	------------	--

AP		INT				
-----------	--	-----	--	--	--	--

AP		ZYDUS PHARMS	1MG/ML	A214655	001	Apr 15, 2022
-----------	--	--------------	---------------	----------------	------------	--------------

DODEX

AP		ACCORD HLTHCARE	1MG/ML	A083022	001	
-----------	--	-----------------	---------------	----------------	------------	--

VIBISON

AP	+	FRESENIUS KABI USA	1MG/ML	A080557	003	
-----------	---	--------------------	---------------	----------------	------------	--

SPRAY, METERED; NASAL

CYANOCOBALAMIN

AB		LUPIN	0.5MG/SPRAY	A210629	001	Jun 30, 2023
-----------	--	-------	--------------------	----------------	------------	--------------

AB	!	PADAGIS ISRAEL	0.5MG/SPRAY	A212458	001	Sep 09, 2020
-----------	---	----------------	--------------------	----------------	------------	--------------

CYCLOBENZAPRINE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

AMRIX

AB	+	TEVA PHARMS INTL	15MG	N021777	001	Feb 01, 2007
-----------	---	------------------	-------------	----------------	------------	--------------

AB	+		30MG	N021777	002	Feb 01, 2007
-----------	---	--	-------------	----------------	------------	--------------

CYCLOBENZAPRINE HYDROCHLORIDE

AB		MACLEODS PHARMS LTD	15MG	A207314	001	Jul 22, 2024
-----------	--	---------------------	-------------	----------------	------------	--------------

AB			30MG	A207314	002	Jul 22, 2024
-----------	--	--	-------------	----------------	------------	--------------

AB		NOVAST LABS	15MG	A214732	001	Aug 27, 2024
-----------	--	-------------	-------------	----------------	------------	--------------

AB			30MG	A214732	002	Aug 27, 2024
-----------	--	--	-------------	----------------	------------	--------------

AB		TWI PHARMS INC	15MG	A091281	001	Jan 31, 2013
-----------	--	----------------	-------------	----------------	------------	--------------

AB			30MG	A091281	002	Jan 31, 2013
-----------	--	--	-------------	----------------	------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

CYCLOBENZAPRINE HYDROCHLORIDE

TABLET; ORAL

CYCLOBENZAPRINE HYDROCHLORIDE

<u>AB</u>	ACTAVIS LABS FL INC	<u>5MG</u>	<u>A071611</u>	<u>002</u>	Feb 03, 2006
<u>AB</u>		<u>7.5MG</u>	<u>A071611</u>	<u>003</u>	Feb 03, 2006
<u>AB</u>		<u>10MG</u>	<u>A071611</u>	<u>001</u>	May 03, 1989
<u>AB</u>	ANDA REPOSITORY	<u>5MG</u>	<u>A073541</u>	<u>002</u>	Apr 06, 2006
<u>AB</u>		<u>10MG</u>	<u>A073541</u>	<u>001</u>	May 23, 1995
<u>AB</u>	AUROBINDO PHARMA	<u>5MG</u>	<u>A078643</u>	<u>001</u>	Sep 26, 2008
<u>AB</u>		<u>10MG</u>	<u>A078643</u>	<u>002</u>	Sep 26, 2008
<u>AB</u>	CHARTWELL RX	<u>5MG</u>	<u>A078218</u>	<u>002</u>	Jun 19, 2015
<u>AB</u>		<u>7.5MG</u>	<u>A078218</u>	<u>003</u>	Nov 03, 2020
<u>AB</u>		<u>10MG</u>	<u>A078218</u>	<u>001</u>	Apr 18, 2008
<u>AB</u>	GRAVITI PHARMS	<u>5MG</u>	<u>A218936</u>	<u>001</u>	Sep 12, 2024
<u>AB</u>		<u>7.5MG</u>	<u>A218936</u>	<u>002</u>	Sep 12, 2024
<u>AB</u>		<u>10MG</u>	<u>A218936</u>	<u>003</u>	Sep 12, 2024
<u>AB</u>	INVAGEN PHARMS	<u>5MG</u>	<u>A090478</u>	<u>001</u>	Jul 23, 2010
<u>AB</u>		<u>10MG</u>	<u>A090478</u>	<u>002</u>	Jul 23, 2010
<u>AB</u>	JUBILANT CADISTA	<u>5MG</u>	<u>A077563</u>	<u>001</u>	Apr 19, 2006
<u>AB</u>		<u>7.5MG</u>	<u>A077563</u>	<u>003</u>	Aug 25, 2017
<u>AB</u>		<u>10MG</u>	<u>A077563</u>	<u>002</u>	Apr 19, 2006
<u>AB</u>	KVK TECH	<u>5MG</u>	<u>A078048</u>	<u>001</u>	Feb 28, 2011
<u>AB</u>		<u>10MG</u>	<u>A078048</u>	<u>002</u>	Feb 28, 2011
<u>AB</u>	OXFORD PHARMS	<u>5MG</u>	<u>A077209</u>	<u>002</u>	Feb 03, 2006
<u>AB</u>		<u>10MG</u>	<u>A077209</u>	<u>001</u>	Oct 04, 2005
<u>AB</u>	PRINSTON INC	<u>5MG</u>	<u>A077797</u>	<u>001</u>	Feb 28, 2007
<u>AB</u>		<u>10MG</u>	<u>A077797</u>	<u>002</u>	Feb 28, 2007
<u>AB</u>	RUBICON	<u>5MG</u>	<u>A208170</u>	<u>001</u>	May 31, 2017
<u>AB</u>		<u>7.5MG</u>	<u>A208170</u>	<u>002</u>	May 31, 2017
<u>AB</u>	!	<u>10MG</u>	<u>A208170</u>	<u>003</u>	May 31, 2017
<u>AB</u>	SUN PHARM INDS LTD	<u>5MG</u>	<u>A078722</u>	<u>001</u>	May 12, 2008
<u>AB</u>		<u>7.5MG</u>	<u>A078722</u>	<u>002</u>	May 12, 2008
<u>AB</u>		<u>10MG</u>	<u>A078722</u>	<u>003</u>	May 12, 2008
<u>AB</u>	UNICHEM	<u>5MG</u>	<u>A213324</u>	<u>001</u>	Jul 06, 2020
<u>AB</u>		<u>7.5MG</u>	<u>A213324</u>	<u>002</u>	Jul 06, 2020
<u>AB</u>		<u>10MG</u>	<u>A213324</u>	<u>003</u>	Jul 05, 2020

CYCLOPENTOLATE HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

AKPENTOLATE

<u>AT</u>	RISING	<u>1%</u>	<u>A040164</u>	<u>001</u>	Jan 13, 1997
<u>AT</u>		<u>2%</u>	<u>A040165</u>	<u>001</u>	Jan 13, 1997

CYCLOGYL

<u>AT</u>	+!	ALCON LABS INC	<u>1%</u>	<u>A084110</u>	<u>001</u>
<u>AT</u>	+!		<u>2%</u>	<u>A084108</u>	<u>001</u>

PENTOLAIR

<u>AT</u>	BAUSCH AND LOMB	<u>1%</u>	<u>A040075</u>	<u>001</u>	Apr 29, 1994
	CYCLOGYL				
	+!	ALCON LABS INC	0.5%	A084109	001

CYCLOPENTOLATE HYDROCHLORIDE; PHENYLEPHRINE HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

CYCLOMYDRIL

!	ALCON LABS INC	0.2%;1%	A084300	001
---	----------------	---------	---------	-----

CYCLOPHOSPHAMIDE

CAPSULE; ORAL

CYCLOPHOSPHAMIDE

<u>AB</u>	ALEMBIC	<u>25MG</u>	<u>A215892</u>	<u>001</u>	Nov 10, 2022	
<u>AB</u>		<u>50MG</u>	<u>A215892</u>	<u>002</u>	Nov 10, 2022	
<u>AB</u>	CIPLA	<u>25MG</u>	<u>A211608</u>	<u>001</u>	Jan 18, 2019	
<u>AB</u>		<u>50MG</u>	<u>A211608</u>	<u>002</u>	Jan 18, 2019	
<u>AB</u>	+	HIKMA	<u>25MG</u>	<u>N203856</u>	<u>001</u>	Sep 16, 2013
<u>AB</u>	+!		<u>50MG</u>	<u>N203856</u>	<u>002</u>	Sep 16, 2013
<u>AB</u>	KANCHAN HLTHCARE	<u>25MG</u>	<u>A209872</u>	<u>001</u>	May 07, 2018	
<u>AB</u>		<u>50MG</u>	<u>A209872</u>	<u>002</u>	May 07, 2018	
<u>AB</u>	NEXTSOURCE	<u>25MG</u>	<u>A218282</u>	<u>001</u>	Dec 03, 2024	
<u>AB</u>		<u>50MG</u>	<u>A218282</u>	<u>002</u>	Dec 03, 2024	
<u>AB</u>	ZYDUS LIFESCIENCES	<u>25MG</u>	<u>A211552</u>	<u>001</u>	Dec 13, 2023	
<u>AB</u>		<u>50MG</u>	<u>A211552</u>	<u>002</u>	Dec 13, 2023	

INJECTABLE; INJECTION

CYCLOPHOSPHAMIDE

<u>AP</u>	AMNEAL	<u>500MG/VIAL</u>	<u>A210046</u>	<u>001</u>	May 25, 2018
<u>AP</u>		<u>1GM/VIAL</u>	<u>A210046</u>	<u>002</u>	May 25, 2018

PRESCRIPTION DRUG PRODUCT LIST

CYCLOPHOSPHAMIDE

INJECTABLE; INJECTION

CYCLOPHOSPHAMIDE

<u>AP</u>			<u>2GM/VIAL</u>	<u>A210046</u>	<u>003</u>	May 25, 2018
<u>AP</u>	!	BAXTER HLTHCARE	<u>500MG/VIAL</u>	<u>A040745</u>	<u>001</u>	May 21, 2008
<u>AP</u>	!		<u>1GM/VIAL</u>	<u>A040745</u>	<u>002</u>	May 21, 2008
<u>AP</u>	!		<u>2GM/VIAL</u>	<u>A040745</u>	<u>003</u>	May 21, 2008
<u>AP</u>		EPIC PHARMA LLC	<u>500MG/VIAL</u>	<u>A218644</u>	<u>001</u>	Dec 19, 2024
<u>AP</u>			<u>1GM/VIAL</u>	<u>A218644</u>	<u>002</u>	Dec 19, 2024
<u>AP</u>			<u>2GM/VIAL</u>	<u>A218644</u>	<u>003</u>	Dec 19, 2024
<u>AP</u>		HAINAN POLY	<u>500MG/VIAL</u>	<u>A218632</u>	<u>001</u>	Dec 26, 2024
<u>AP</u>			<u>1GM/VIAL</u>	<u>A218632</u>	<u>002</u>	Dec 26, 2024
<u>AP</u>			<u>2GM/VIAL</u>	<u>A218632</u>	<u>003</u>	Dec 26, 2024
<u>AP</u>		HENGRUI PHARMA	<u>500MG/VIAL</u>	<u>A204555</u>	<u>001</u>	Oct 31, 2014
<u>AP</u>			<u>1GM/VIAL</u>	<u>A204555</u>	<u>002</u>	Oct 31, 2014
<u>AP</u>			<u>2GM/VIAL</u>	<u>A204555</u>	<u>003</u>	Oct 31, 2014
<u>AP</u>		HIKMA	<u>500MG/VIAL</u>	<u>A216958</u>	<u>001</u>	Dec 18, 2023
<u>AP</u>			<u>1GM/VIAL</u>	<u>A216958</u>	<u>002</u>	Dec 18, 2023
<u>AP</u>			<u>2GM/VIAL</u>	<u>A216958</u>	<u>003</u>	Dec 18, 2023
<u>AP</u>		SAGENT PHARMS INC	<u>500MG/VIAL</u>	<u>A214529</u>	<u>001</u>	Jul 17, 2023
<u>AP</u>			<u>1GM/VIAL</u>	<u>A214529</u>	<u>002</u>	Jul 17, 2023
<u>AP</u>			<u>2GM/VIAL</u>	<u>A214529</u>	<u>003</u>	Jul 17, 2023
<u>AP</u>		SUNNY	<u>500MG/VIAL</u>	<u>A215089</u>	<u>001</u>	Oct 26, 2023
<u>AP</u>			<u>1GM/VIAL</u>	<u>A215089</u>	<u>002</u>	Oct 26, 2023
<u>AP</u>			<u>2GM/VIAL</u>	<u>A215089</u>	<u>003</u>	Oct 26, 2023
<u>AP</u>		XGEN PHARMS	<u>500MG/VIAL</u>	<u>A211757</u>	<u>001</u>	Oct 18, 2022
<u>AP</u>			<u>1GM/VIAL</u>	<u>A211757</u>	<u>002</u>	Oct 18, 2022
<u>AP</u>			<u>2GM/VIAL</u>	<u>A211757</u>	<u>003</u>	Oct 18, 2022

SOLUTION; INTRAVENOUS

CYCLOPHOSPHAMIDE

<u>AP</u>	+	DR REDDYS	<u>500MG/2.5ML (200MG/ML)</u>	<u>N212501</u>	<u>001</u>	Jul 30, 2020
<u>AP</u>	+		<u>1GM/5ML (200MG/ML)</u>	<u>N212501</u>	<u>002</u>	Jul 30, 2020
<u>AP</u>		NEXUS	<u>500MG/2.5ML (200MG/ML)</u>	<u>A216783</u>	<u>001</u>	Oct 29, 2024
<u>AP</u>			<u>1GM/5ML (200MG/ML)</u>	<u>A216783</u>	<u>002</u>	Oct 29, 2024
	+	AVYXA HOLDINGS	500MG/ML (500MG/ML)	N210852	001	Jun 07, 2023
	+		1GM/2ML (500MG/ML)	N210852	002	Jun 07, 2023
	+		2GM/4ML (500MG/ML)	N210852	003	Jun 07, 2023
	+	BAXTER HLTHCARE CORP	500MG/2.5ML (200MG/ML)	N217651	001	Jun 28, 2023
	+		1GM/5ML (200MG/ML)	N217651	002	Jun 28, 2023
	+	DR REDDYS	2GM/10ML (200MG/ML)	N212501	003	Nov 19, 2021
	+	EUGIA PHARMA SPECLTS	500MG/2.5ML (200MG/ML)	N210735	001	Aug 25, 2021
	+		1GM/5ML (200MG/ML)	N210735	002	Aug 25, 2021
	+		2GM/10ML (200MG/ML)	N210735	003	Nov 20, 2023
	+	SANDOZ	500MG/5ML (100MG/ML)	N217150	001	Sep 12, 2023
	+		1GM/10ML (100MG/ML)	N217150	002	Sep 12, 2023
	+		2GM/20ML (100MG/ML)	N217150	003	Sep 12, 2023

TABLET; ORAL

CYTOXAN

+	BAXTER HLTHCARE	25MG	N012141	002
+		50MG	N012141	001

CYCLOSERINE

CAPSULE; ORAL

SEROMYCIN

!	SANALUZ	250MG	A060593	001
---	---------	-------	---------	-----

CYCLOSPORINE

CAPSULE; ORAL

CYCLOSPORINE

<u>AB1</u>		APOTEX	<u>25MG</u>	<u>A210721</u>	<u>001</u>	Jul 10, 2019
<u>AB1</u>			<u>50MG</u>	<u>A210721</u>	<u>002</u>	Jul 10, 2019
<u>AB1</u>			<u>100MG</u>	<u>A210721</u>	<u>003</u>	Jul 10, 2019
<u>AB1</u>		DR REDDYS LABS SA	<u>25MG</u>	<u>A065044</u>	<u>002</u>	Dec 20, 2000
<u>AB1</u>			<u>100MG</u>	<u>A065044</u>	<u>001</u>	Dec 20, 2000
<u>AB1</u>		IVAX SUB TEVA PHARMS	<u>25MG</u>	<u>A065110</u>	<u>003</u>	Mar 29, 2005
<u>AB1</u>			<u>50MG</u>	<u>A065110</u>	<u>001</u>	Mar 29, 2005
<u>AB1</u>			<u>100MG</u>	<u>A065110</u>	<u>002</u>	Mar 29, 2005
<u>AB1</u>		SANDOZ	<u>25MG</u>	<u>A065017</u>	<u>002</u>	Jan 13, 2000
<u>AB1</u>			<u>100MG</u>	<u>A065017</u>	<u>001</u>	Jan 13, 2000
<u>AB1</u>		STRIDES SOFTGELS	<u>25MG</u>	<u>A216046</u>	<u>001</u>	Aug 02, 2022
<u>AB1</u>			<u>50MG</u>	<u>A216046</u>	<u>002</u>	Aug 02, 2022

PRESCRIPTION DRUG PRODUCT LIST

CYCLOSPORINE

CAPSULE; ORAL

CYCLOSPORINE**AB1** **100MG** **A216046 003** Aug 02, 2022GENGRAF**AB1** ABBVIE **25MG** **A065003 001** May 12, 2000**AB1** **100MG** **A065003 003** May 12, 2000NEORAL**AB1** + NOVARTIS **25MG** **N050715 001** Jul 14, 1995**AB1** +! **100MG** **N050715 002** Jul 14, 1995CYCLOSPORINE**AB2** APOTEX **25MG** **A065040 001** May 09, 2002**AB2** **100MG** **A065040 002** May 09, 2002SANDIMMUNE**AB2** + NOVARTIS **25MG** **N050625 001** Mar 02, 1990**AB2** +! **100MG** **N050625 002** Mar 02, 1990BX + **50MG** **N050625 003** Nov 23, 1992

EMULSION; OPHTHALMIC

CYCLOSPORINE**AB** APOTEX **0.05%** **A207606 001** Jan 12, 2023**AB** DEVA HOLDING AS **0.05%** **A209811 001** May 21, 2024**AB** MYLAN **0.05%** **A205894 001** Feb 02, 2022**AB** SAPTALIS PHARMS **0.05%** **A211943 001** Jul 05, 2024**AB** TEVA PHARMS USA INC **0.05%** **A203880 001** Dec 14, 2023RESTASIS**AB** +! ABBVIE **0.05%** **N050790 001** Dec 23, 2002

RESTASIS MULTIDOSE

+! ABBVIE **0.05%** **N050790 002** Oct 27, 2016

VERKAZIA

+! HARROW EYE **0.1%** **N214965 001** Jun 23, 2021

INJECTABLE; INJECTION

CYCLOSPORINE**AP** HIKMA **50MG/ML** **A065004 001** Oct 29, 1999**AP** PADAGIS US **50MG/ML** **A065151 001** Oct 07, 2003SANDIMMUNE**AP** +! NOVARTIS **50MG/ML** **N050573 001** Nov 14, 1983

SOLUTION; OPHTHALMIC

CEQUA

+! SUN PHARM **0.09%** **N210913 001** Aug 14, 2018

VEVYE

+! HARROW EYE **0.1%** **N217469 001** May 30, 2023

SOLUTION; ORAL

CYCLOSPORINE**AB1** ABBVIE **100MG/ML** **A065025 001** Mar 03, 2000**AB1** IVAX SUB TEVA **100MG/ML** **A065078 001** Mar 25, 2005

PHARMS

NEORAL**AB1** +! NOVARTIS **100MG/ML** **N050716 001** Jul 14, 1995

SANDIMMUNE

+! NOVARTIS **100MG/ML** **N050574 001** Nov 14, 1983CYPROHEPTADINE HYDROCHLORIDE

SYRUP; ORAL

CYPROHEPTADINE HYDROCHLORIDE**AA** CHARTWELL MOLECULAR **2MG/5ML** **A203191 001** Jul 13, 2017**AA** ELYSIUM **2MG/5ML** **A209108 001** Oct 16, 2018**AA** QUAGEN **2MG/5ML** **A212423 001** May 22, 2019**AA** ! RISING **2MG/5ML** **A040668 001** Jun 28, 2006

TABLET; ORAL

CYPROHEPTADINE HYDROCHLORIDE**AA** APPCO **4MG** **A206553 001** Nov 29, 2016**AA** BEXIMCO PHARMS USA **4MG** **A206676 001** Apr 12, 2019**AA** CHARTWELL RX **4MG** **A088212 001** May 26, 1983**AA** KENTON **4MG** **A040644 001** May 30, 2006**AA** MOUNTAIN **4MG** **A040537 001** Sep 30, 2003**AA** NOVAST LABS **4MG** **A205087 001** Sep 23, 2015**AA** QUAGEN **4MG** **A212491 001** Feb 24, 2021**AA** RISING **4MG** **A207555 001** Jan 31, 2017**AA** STRIDES PHARMA **4MG** **A209172 001** Apr 11, 2018**AA** ! ZYDUS PHARMS **4MG** **A208938 001** May 19, 2017

PRESCRIPTION DRUG PRODUCT LIST

CYSTEAMINE BITARTRATE

CAPSULE; ORAL

CYSTAGON

+ MYLAN

EQ 50MG BASE

N020392 001 Aug 15, 1994

+!

EQ 150MG BASE

N020392 002 Aug 15, 1994

CAPSULE, DELAYED RELEASE; ORAL

PROCYSBI

+ HORIZON

EQ 25MG BASE

N203389 001 Apr 30, 2013

+!

EQ 75MG BASE

N203389 002 Apr 30, 2013

GRANULE, DELAYED RELEASE; ORAL

PROCYSBI

+ HORIZON

EQ 75MG BASE/PACKET

N213491 001 Feb 14, 2020

+!

EQ 300MG BASE/PACKET

N213491 002 Feb 14, 2020

CYSTEAMINE HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

CYSTADROPS

+! RECORDATI RARE

EQ 0.37% BASE

N211302 001 Aug 19, 2020

CYSTARAN

+! LEADIANT BIOSCI INC

EQ 0.44% BASE

N200740 001 Oct 02, 2012

CYSTEINE HYDROCHLORIDE

SOLUTION; INTRAVENOUS

ELCYS

+! EXELA PHARMA

500MG/10ML (50MG/ML)

N210660 001 Apr 16, 2019

CYTARABINE

INJECTABLE; INJECTION

CYTARABINE**AP** ! FRESENIUS KABI USA**100MG/ML****A076512 001** Jan 15, 2004**AP** GLAND PHARMA LTD**100MG/VIAL****A211937 001** Dec 23, 2019**AP****2GM/VIAL****A211938 001** Dec 23, 2019**AP** HIKMA**100MG/VIAL****A071471 001** Aug 02, 1989**AP** ! HOSPIRA**20MG/ML****A071868 001** Jun 04, 1990**AP** !**20MG/ML****A072168 001** Aug 31, 1990**AP** !**20MG/ML****A072945 001** Feb 28, 1994**AP****100MG/ML****A075383 001** Nov 22, 1999**AP** MEITHEAL**20MG/ML****A208485 001** Feb 28, 2022**AP****100MG/ML****A205696 001** Jul 17, 2018**AP** RISING**20MG/ML****A200915 001** Dec 13, 2011**AP****100MG/ML****A201784 001** Jan 30, 2012**AP** ! WEST-WARD PHARMS**2GM/VIAL****A074245 002** Aug 31, 1994

! HIKMA

500MG/VIAL

A071472 001 Aug 02, 1989

! WEST-WARD PHARMS

1GM/VIAL

A074245 001 Aug 31, 1994

! HIKMA

500MG/VIAL

A071472 001 Aug 02, 1989

! WEST-WARD PHARMS

1GM/VIAL

A074245 001 Aug 31, 1994

! HIKMA

500MG/VIAL

A071472 001 Aug 02, 1989

! WEST-WARD PHARMS

1GM/VIAL

A074245 001 Aug 31, 1994

! HIKMA

500MG/VIAL

A071472 001 Aug 02, 1989

! WEST-WARD PHARMS

1GM/VIAL

A074245 001 Aug 31, 1994

! HIKMA

500MG/VIAL

A071472 001 Aug 02, 1989

! WEST-WARD PHARMS

1GM/VIAL

A074245 001 Aug 31, 1994

! HIKMA

500MG/VIAL

A071472 001 Aug 02, 1989

! WEST-WARD PHARMS

1GM/VIAL

A074245 001 Aug 31, 1994

! HIKMA

500MG/VIAL

A071472 001 Aug 02, 1989

! WEST-WARD PHARMS

1GM/VIAL

A074245 001 Aug 31, 1994

! HIKMA

500MG/VIAL

A071472 001 Aug 02, 1989

! WEST-WARD PHARMS

1GM/VIAL

A074245 001 Aug 31, 1994

! HIKMA

500MG/VIAL

A071472 001 Aug 02, 1989

! WEST-WARD PHARMS

1GM/VIAL

A074245 001 Aug 31, 1994

! HIKMA

500MG/VIAL

A071472 001 Aug 02, 1989

! WEST-WARD PHARMS

1GM/VIAL

A074245 001 Aug 31, 1994

! HIKMA

500MG/VIAL

A071472 001 Aug 02, 1989

! WEST-WARD PHARMS

1GM/VIAL

A074245 001 Aug 31, 1994

! HIKMA

500MG/VIAL

A071472 001 Aug 02, 1989

! WEST-WARD PHARMS

1GM/VIAL

A074245 001 Aug 31, 2020

! HIKMA

500MG/VIAL

A207961 003 Aug 13, 2024

! WEST-WARD PHARMS

1GM/VIAL

A207961 002 May 06, 2020

! HIKMA

500MG/VIAL

A213879 001 May 22, 2024

! WEST-WARD PHARMS

1GM/VIAL

A213879 003 Aug 12, 2024

! HIKMA

500MG/VIAL

A213879 002 May 22, 2024

! WEST-WARD PHARMS

1GM/VIAL

A213879 002 May 22, 2024

! HIKMA

500MG/VIAL

A213879 002 May 22, 2024

! WEST-WARD PHARMS

1GM/VIAL

A213879 002 May 22, 2024

! HIKMA

500MG/VIAL

A213879 002 May 22, 2024

! WEST-WARD PHARMS

1GM/VIAL

A213879 002 May 22, 2024

! HIKMA

500MG/VIAL

A213879 002 May 22, 2024

! WEST-WARD PHARMS

1GM/VIAL

A213879 002 May 22, 2024

PRADAXA**AB** + BOEHRINGER**EQ 75MG BASE****N022512 001** Oct 19, 2010

PRESCRIPTION DRUG PRODUCT LIST

DABIGATRAN ETEXILATE MESYLATE

CAPSULE; ORAL

PRADAXA

INGELHEIM

AB	+		<u>EQ 110MG BASE</u>	<u>N022512 003</u>	Nov 20, 2015
AB	+	!	<u>EQ 150MG BASE</u>	<u>N022512 002</u>	Oct 19, 2010

PELLETS; ORAL

PRADAXA+ BOEHRINGER
INGELHEIM

			EQ 20MG BASE/PACKET	N214358 001	Jun 21, 2021
			EQ 30MG BASE/PACKET	N214358 002	Jun 21, 2021
			EQ 40MG BASE/PACKET	N214358 003	Jun 21, 2021
			EQ 50MG BASE/PACKET	N214358 004	Jun 21, 2021
			EQ 110MG BASE/PACKET	N214358 005	Jun 21, 2021
		+	EQ 150MG BASE/PACKET	N214358 006	Jun 21, 2021

DABRAFENIB MESYLATE

CAPSULE; ORAL

TAFINLAR

+ NOVARTIS

EQ 50MG BASE N202806 001 May 29, 2013

+!

EQ 75MG BASE N202806 002 May 29, 2013

TABLET, FOR SUSPENSION; ORAL

TAFINLAR

+! NOVARTIS

EQ 10MG BASE N217514 001 Mar 16, 2023

DACARBAZINE

INJECTABLE; INJECTION

DACARBAZINE

AP	!	FRESENIUS KABI USA	<u>200MG/VIAL</u>	<u>A075371 002</u>	Aug 27, 1999
AP		HIKMA	<u>200MG/VIAL</u>	<u>A075812 001</u>	Jun 15, 2001
AP			<u>500MG/VIAL</u>	<u>A075812 002</u>	Oct 31, 2002
AP		HOSPIRA	<u>200MG/VIAL</u>	<u>A075940 001</u>	Oct 18, 2001
AP		MEITHEAL	<u>200MG/VIAL</u>	<u>A075259 002</u>	Aug 27, 1998
AP	!		<u>500MG/VIAL</u>	<u>A075259 001</u>	Sep 22, 2000
	!	FRESENIUS KABI USA	100MG/VIAL	A075371 001	Aug 27, 1999

DACOMITINIB

TABLET; ORAL

VIZIMPRO

+ PFIZER

15MG N211288 001 Sep 27, 2018

+

30MG N211288 002 Sep 27, 2018

+!

45MG N211288 003 Sep 27, 2018

DACTINOMYCIN

INJECTABLE; INJECTION

DACTINOMYCIN

AP	!	EUGIA PHARMA	<u>0.5MG/VIAL</u>	<u>A203385 001</u>	Nov 09, 2017
AP		HISUN PHARM	<u>0.5MG/VIAL</u>	<u>A207232 001</u>	Jul 16, 2019
		HANGZHOU			
AP		MEITHEAL	<u>0.5MG/VIAL</u>	<u>A213463 001</u>	Nov 13, 2020
AP		XGEN PHARMS	<u>0.5MG/VIAL</u>	<u>A203999 001</u>	May 20, 2019

DALBAVANCIN HYDROCHLORIDE

POWDER; INTRAVENOUS

DALVANCE

+! ABBVIE

EQ 500MG BASE/VIAL N021883 001 May 23, 2014

DALFAMPRIDINE

TABLET, EXTENDED RELEASE; ORAL

AMPYRA

AB	+	!	MERZ PHARMS	<u>10MG</u>	<u>N022250 001</u>	Jan 22, 2010
-----------	---	---	-------------	--------------------	---------------------------	--------------

DALFAMPRIDINE

AB			ACCORD HLTHCARE	<u>10MG</u>	<u>A206863 001</u>	Jul 11, 2018
AB			ACTAVIS LABS FL INC	<u>10MG</u>	<u>A206836 001</u>	Jan 23, 2017
AB			ALKEM LABS LTD	<u>10MG</u>	<u>A206765 001</u>	Jul 30, 2018
AB			AUROBINDO PHARMA	<u>10MG</u>	<u>A206811 001</u>	Jan 23, 2017
AB			MICRO LABS	<u>10MG</u>	<u>A210158 001</u>	Mar 11, 2019
AB			SUN PHARM	<u>10MG</u>	<u>A208292 001</u>	May 21, 2019

DALTEPARIN SODIUM

INJECTABLE; SUBCUTANEOUS

FRAGMIN

+ PFIZER

2,500IU/0.2ML (12,500IU/ML) N020287 001 Dec 22, 1994

+

5,000IU/0.2ML (25,000IU/ML) N020287 003 Mar 18, 1996

+

7,500IU/0.3ML (25,000IU/ML) N020287 005 Apr 04, 2002

+

10,000IU/ML (10,000IU/ML) N020287 004 Jan 30, 1998

PRESCRIPTION DRUG PRODUCT LIST

DALTEPARIN SODIUM

INJECTABLE; SUBCUTANEOUS

FRAGMIN

+	10,000IU/4ML (2,500IU/ML)	N020287 012	Mar 15, 2022
+	12,500IU/0.5ML (25,000IU/ML)	N020287 009	May 01, 2007
+	15,000IU/0.6ML (25,000IU/ML)	N020287 010	May 01, 2007
+	18,000IU/0.72ML (25,000IU/ML)	N020287 011	May 01, 2007
+	95,000IU/3.8ML (25,000IU/ML)	N020287 006	Apr 04, 2002

DANAZOL

CAPSULE; ORAL

DANAZOL

<u>AB</u>	BARR	<u>50MG</u>	<u>A074582 003</u>	May 29, 1998
<u>AB</u>		<u>100MG</u>	<u>A074582 002</u>	May 29, 1998
<u>AB</u>	!	<u>200MG</u>	<u>A074582 001</u>	Aug 09, 1996
<u>AB</u>	LANNETT CO INC	<u>50MG</u>	<u>A077246 002</u>	Apr 19, 2007
<u>AB</u>		<u>100MG</u>	<u>A077246 003</u>	Apr 19, 2007
<u>AB</u>		<u>200MG</u>	<u>A077246 001</u>	Sep 28, 2005

DANICOPAN

TABLET; ORAL

VOYDEYA

+	ALEXION PHARMS INC	50MG	N218037 001	Mar 29, 2024
+		100MG	N218037 002	Mar 29, 2024

DANTROLENE SODIUM

CAPSULE; ORAL

DANTRIUM

<u>AB</u>	+	ENDO OPERATIONS	<u>25MG</u>	<u>N017443 001</u>
<u>AB</u>	+		<u>50MG</u>	<u>N017443 003</u>
<u>AB</u>	+		<u>100MG</u>	<u>N017443 002</u>

DANTROLENE SODIUM

<u>AB</u>	ELITE LABS INC	<u>25MG</u>	<u>A076686 001</u>	Oct 24, 2005
<u>AB</u>		<u>50MG</u>	<u>A076686 002</u>	Oct 24, 2005
<u>AB</u>		<u>100MG</u>	<u>A076686 003</u>	Oct 24, 2005
<u>AB</u>	IMPAX LABS	<u>25MG</u>	<u>A076856 001</u>	Mar 01, 2005
<u>AB</u>		<u>50MG</u>	<u>A076856 002</u>	Mar 01, 2005
<u>AB</u>		<u>100MG</u>	<u>A076856 003</u>	Mar 01, 2005

FOR SUSPENSION; INTRAVENOUS

RYANODEX

+	EAGLE PHARMS	250MG/VIAL	N205579 001	Jul 22, 2014
---	--------------	------------	-------------	--------------

INJECTABLE; INJECTION

DANTRIUM

<u>AP</u>	+	ENDO OPERATIONS	<u>20MG/VIAL</u>	<u>N018264 001</u>
-----------	---	-----------------	------------------	--------------------

DANTROLENE SODIUM

<u>AP</u>	HIKMA	<u>20MG/VIAL</u>	<u>A204762 001</u>	Jun 19, 2017
-----------	-------	------------------	--------------------	--------------

REVONTO

<u>AP</u>	USWM	<u>20MG/VIAL</u>	<u>A078378 001</u>	Jul 24, 2007
-----------	------	------------------	--------------------	--------------

DAPAGLIFLOZIN

TABLET; ORAL

FARXIGA

+	ASTRAZENECA AB	5MG	N202293 001	Jan 08, 2014
+		10MG	N202293 002	Jan 08, 2014

DAPAGLIFLOZIN; METFORMIN HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL

XIGDUO XR

+	ASTRAZENECA AB	2.5MG; 1GM	N205649 005	Jul 28, 2017
+		5MG; 500MG	N205649 001	Oct 29, 2014
+		5MG; 1GM	N205649 002	Oct 29, 2014
+		10MG; 500MG	N205649 003	Oct 29, 2014
+		10MG; 1GM	N205649 004	Oct 29, 2014

DAPAGLIFLOZIN; SAXAGLIPTIN HYDROCHLORIDE

TABLET; ORAL

QTERN

+	ASTRAZENECA AB	5MG; EQ 5MG BASE	N209091 002	May 02, 2019
+		10MG; EQ 5MG BASE	N209091 001	Feb 27, 2017

PRESCRIPTION DRUG PRODUCT LIST

DAPSONE

GEL; TOPICAL

ACZONE

AB	+ !	ABBVIE	5%	N021794 001	Jul 07, 2005
AB	+ !	ALMIRALL	7.5%	N207154 001	Feb 24, 2016

DAPSONE

AB		ALEMBIC	7.5%	A215718 001	Oct 27, 2023
AB		AMNEAL	5%	A209890 001	Oct 19, 2023
AB			7.5%	A212701 001	May 31, 2023
AB		COSETTE	5%	A210178 001	Mar 31, 2022
AB		ENCUBE	5%	A212383 001	Oct 13, 2023
AB		MYLAN	7.5%	A213847 001	Feb 04, 2022
AB		TARO	5%	A209506 001	Oct 16, 2017
AB			7.5%	A210191 001	Jun 26, 2019
AB		TORRENT	7.5%	A214722 001	Feb 10, 2022
AB		TRUPHARMA	5%	A213907 001	Jun 06, 2023
AB			7.5%	A213636 001	Aug 26, 2024
AB		ZYDUS LIFESCIENCES	7.5%	A214019 001	May 08, 2024

TABLET; ORAL

DAPSONE

AB		ACTAVIS LLC	25MG	A204380 001	Mar 23, 2017
AB			100MG	A204380 002	Mar 23, 2017
AB	+	EVEREST LIFE SCI	25MG	A086841 001	
AB	+		100MG	A086842 001	
AB		NOSTRUM LABS INC	25MG	A203887 001	May 06, 2016
AB			100MG	A203887 002	May 06, 2016
AB		NOVITIUM PHARMA	25MG	A206505 001	Dec 01, 2016
AB	!		100MG	A206505 002	Dec 01, 2016
AB		RISING	25MG	A207165 002	Jul 20, 2023
AB			100MG	A207165 001	May 08, 2019

DAPTOMYCIN

POWDER; INTRAVENOUS

DAPTOMYCIN

AP		ACCORD HLTHCARE	350MG/VIAL	A212667 001	Jul 12, 2019
AP			500MG/VIAL	A211961 001	Jun 24, 2019
AP		ASPIRO	350MG/VIAL	A216445 001	Dec 23, 2022
AP			500MG/VIAL	A216413 001	Sep 23, 2022
AP		BE PHARMS	350MG/VIAL	A213425 001	Aug 20, 2020
AP			500MG/VIAL	A212513 001	Jun 26, 2019
AP		BIOCON PHARMA	500MG/VIAL	A217215 001	Aug 29, 2024
AP		DR REDDYS	350MG/VIAL	A211403 001	Aug 31, 2020
AP			500MG/VIAL	A208375 001	May 01, 2019
AP		EUGIA PHARMA	500MG/VIAL	A213171 001	Sep 02, 2021
AP		FRESENIUS KABI USA	350MG/VIAL	A213396 001	Aug 01, 2024
AP			500MG/VIAL	A206077 001	Apr 11, 2018
AP		HAINAN POLY PHARM	500MG/VIAL	A215890 001	Aug 18, 2022
AP		HANGZHOU ZHONGMEI	500MG/VIAL	A215215 001	Nov 26, 2021
AP		HENGRUI PHARMA	500MG/VIAL	A212022 001	Aug 22, 2019
AP		HISUN PHARM	500MG/VIAL	A212250 001	Apr 21, 2021
AP		HANGZHOU			
AP		MEITHEAL	350MG/VIAL	A213786 001	Jun 29, 2021
AP			500MG/VIAL	A213623 001	Jun 29, 2021
AP		MYLAN	500MG/VIAL	A213966 001	Aug 07, 2023
AP		MYLAN LABS LTD	500MG/VIAL	A205037 001	Jun 05, 2018
AP	!	QILU PHARM HAINAN	500MG/VIAL	A215316 001	Aug 24, 2021
AP	+ !	SAGENT PHARMS INC	350MG/VIAL	N208385 001	Sep 12, 2017
AP			500MG/VIAL	A207104 001	Nov 15, 2019
AP		TEVA PHARMS USA	500MG/VIAL	A091039 001	Mar 25, 2016
AP		XELLIA PHARMS APS	500MG/VIAL	A206005 001	Jun 15, 2016
	+ !	HIKMA	350MG/VIAL	N209949 001	Oct 20, 2017
	+ !		350MG/VIAL	N217415 001	Jan 30, 2023
	+ !		500MG/VIAL	N217415 002	Jan 30, 2023
	+ !	HOSPIRA	350MG/VIAL	N210282 001	Jun 21, 2021
	+ !		500MG/VIAL	N210282 002	Jun 21, 2021
	+ !	MAIA PHARMS INC	350MG/VIAL	N217630 001	Nov 21, 2024
	+ !		500MG/VIAL	N217630 002	Nov 21, 2024

SOLUTION; INTRAVENOUS

DAPTOMYCIN IN 0.9% SODIUM CHLORIDE

+ !	BAXTER HLTHCARE	350MG/50ML (7MG/ML)	N213645 002	Feb 27, 2023
+ !	CORP			
+ !		500MG/50ML (10MG/ML)	N213645 003	Feb 27, 2023
+ !		700MG/100ML (7MG/ML)	N213645 004	Feb 27, 2023

PRESCRIPTION DRUG PRODUCT LIST

DAPTOMYCIN

SOLUTION;INTRAVENOUS

DAPTOMYCIN IN 0.9% SODIUM CHLORIDE

+! 1GM/100ML (10MG/ML) N213645 005 Feb 27, 2023

DARIDOREXANT HYDROCHLORIDE

TABLET;ORAL

QUVIVIQ

+ IDORSIA EQ 25MG BASE N214985 001 Apr 07, 2022

+! EQ 50MG BASE N214985 002 Apr 07, 2022

DARIFENACIN HYDROBROMIDE

TABLET, EXTENDED RELEASE;ORAL

DARIFENACIN**AB** MACLEODS PHARMS LTD **EQ 7.5MG BASE** **A207302 001** Jul 28, 2017**AB** ! **EQ 15MG BASE** **A207302 002** Jul 28, 2017DARIFENACIN HYDROBROMIDE**AB** ALEMBIC **EQ 7.5MG BASE** **A207681 001** Dec 08, 2017**AB** **EQ 15MG BASE** **A207681 002** Dec 08, 2017**AB** AUROBINDO PHARMA **EQ 7.5MG BASE** **A206743 001** Sep 19, 2016**AB** **EQ 15MG BASE** **A206743 002** Sep 19, 2016**AB** CIPLA **EQ 7.5MG BASE** **A207664 001** Sep 01, 2016**AB** **EQ 15MG BASE** **A207664 002** Sep 01, 2016**AB** POLYGEN PHARMS **EQ 7.5MG BASE** **A211045 001** Jan 06, 2020**AB** **EQ 15MG BASE** **A211045 002** Jan 06, 2020**AB** TORRENT **EQ 7.5MG BASE** **A205209 001** Nov 17, 2016**AB** **EQ 15MG BASE** **A205209 002** Nov 17, 2016DAROLUTAMIDE

TABLET;ORAL

NUBEQA

+! BAYER HEALTHCARE 300MG N212099 001 Jul 30, 2019

DARUNAVIR

SUSPENSION;ORAL

PREZISTA

+! JANSSEN PRODS 100MG/ML N202895 001 Dec 16, 2011

TABLET;ORAL

DARUNAVIR**AB** AMNEAL **600MG** **A212493 001** Dec 08, 2023**AB** **800MG** **A212493 002** Dec 08, 2023**AB** AUROBINDO PHARMA **600MG** **A210677 001** Nov 28, 2023

LTD

AB **800MG** **A210677 002** Nov 28, 2023**AB** CIPLA **600MG** **A206288 001** Nov 28, 2023**AB** **800MG** **A206288 002** Nov 28, 2023**AB** DR REDDYS **600MG** **A211578 001** Nov 28, 2023**AB** **800MG** **A211578 002** Nov 28, 2023**AB** HETERO LABS LTD III **600MG** **A202083 002** Sep 14, 2023**AB** LUPIN LTD **600MG** **A202073 001** Sep 29, 2022**AB** **800MG** **A202073 002** Sep 29, 2022**AB** MSN **600MG** **A215389 001** Nov 28, 2023**AB** **800MG** **A215389 002** Nov 28, 2023**AB** TEVA PHARMS USA **600MG** **A202118 001** Nov 21, 2017**AB** ZYDUS LIFESCIENCES **600MG** **A214085 001** Dec 13, 2023**AB** **800MG** **A214085 002** Dec 13, 2023PREZISTA**AB** + JANSSEN PRODS **600MG** **N021976 002** Feb 25, 2008**AB** +! **800MG** **N021976 006** Nov 09, 2012

DARUNAVIR

HETERO LABS LTD III 400MG A202083 001 Sep 14, 2023

PREZISTA

+ JANSSEN PRODS 75MG N021976 004 Dec 18, 2008

+ 150MG N021976 005 Dec 18, 2008

DASATINIB

TABLET;ORAL

DASATINIB**AB** APOTEX **20MG** **A202103 001** Jun 10, 2016**AB** **50MG** **A202103 002** Jun 10, 2016**AB** **70MG** **A202103 003** Jun 10, 2016**AB** **80MG** **A203180 001** Nov 23, 2021**AB** **100MG** **A202103 004** Jun 10, 2016**AB** **140MG** **A203180 002** Nov 23, 2021

PRESCRIPTION DRUG PRODUCT LIST

DASATINIB

TABLET; ORAL

SPRYCEL

<u>AB</u>	+	BRISTOL MYERS SQUIBB	<u>20MG</u>	<u>N021986 001</u>	Jun 28, 2006
<u>AB</u>	+		<u>50MG</u>	<u>N021986 002</u>	Jun 28, 2006
<u>AB</u>	+		<u>70MG</u>	<u>N021986 003</u>	Jun 28, 2006
<u>AB</u>	+		<u>80MG</u>	<u>N021986 005</u>	Oct 28, 2010
<u>AB</u>	+		<u>100MG</u>	<u>N021986 004</u>	May 30, 2008
<u>AB</u>	+		<u>140MG</u>	<u>N021986 006</u>	Oct 28, 2010
<u>PHYRAGO</u>					
	+	NANOCOPOEIA	20MG	N216099 001	Dec 05, 2023
	+		50MG	N216099 002	Dec 05, 2023
	+		70MG	N216099 003	Dec 05, 2023
	+		80MG	N216099 004	Dec 05, 2023
	+		100MG	N216099 005	Dec 05, 2023
	+		140MG	N216099 006	Dec 05, 2023

DASIGLUCAGON HYDROCHLORIDE

SOLUTION; SUBCUTANEOUS

ZEGALOGUE

	+	ZEALAND PHARMA	EQ 0.6MG BASE/0.6ML (EQ 0.6MG BASE/0.6ML)	N214231 001	Mar 22, 2021
<u>ZEGALOGUE (AUTOINJECTOR)</u>					
	+	ZEALAND PHARMA	EQ 0.6MG BASE/0.6ML (EQ 0.6MG BASE/0.6ML)	N214231 002	Mar 22, 2021

DAUNORUBICIN HYDROCHLORIDE

INJECTABLE; INJECTION

CERUBIDINE

<u>AP</u>	!	HIKMA	<u>EQ 20MG BASE/VIAL</u>	<u>A064103 001</u>	Feb 03, 1995
<u>DAUNORUBICIN HYDROCHLORIDE</u>					
<u>AP</u>		FRESENIUS KABI USA	<u>EQ 20MG BASE/VIAL</u>	<u>A065000 001</u>	May 25, 1999
<u>AP</u>	+	HIKMA	<u>EQ 5MG BASE/ML</u>	<u>N050731 001</u>	Jan 30, 1998
<u>AP</u>		HISUN PHARM HANGZHOU	<u>EQ 5MG BASE/ML</u>	<u>A208759 001</u>	Apr 12, 2019
<u>AP</u>		MEITHEAL FRESENIUS KABI USA	<u>EQ 5MG BASE/ML</u>	<u>A065035 001</u>	Jan 24, 2000
			EQ 5MG BASE/VIAL	A065034 001	Nov 20, 2001

DECITABINE

INJECTABLE; INTRAVENOUS

DECITABINE

<u>AP</u>	!	ACCORD HLTHCARE	<u>50MG/VIAL</u>	<u>A203475 001</u>	Feb 27, 2017
<u>AP</u>		CHEMI SPA	<u>50MG/VIAL</u>	<u>A206033 001</u>	Sep 22, 2017
<u>AP</u>		DR REDDYS	<u>50MG/VIAL</u>	<u>A203131 001</u>	Jul 11, 2013
<u>AP</u>		EUGIA PHARMA	<u>50MG/VIAL</u>	<u>A214569 001</u>	Sep 20, 2021
<u>AP</u>		GLAND	<u>50MG/VIAL</u>	<u>A205539 001</u>	Nov 23, 2020
<u>AP</u>		HETERO LABS LTD VI	<u>50MG/VIAL</u>	<u>A215355 001</u>	May 10, 2024
<u>AP</u>		JIANGSU HANSOH PHARM	<u>50MG/VIAL</u>	<u>A213472 001</u>	Apr 15, 2022
<u>AP</u>		LUPIN LTD	<u>50MG/VIAL</u>	<u>A210756 001</u>	Nov 09, 2018
<u>AP</u>		MEITHEAL	<u>50MG/VIAL</u>	<u>A212959 001</u>	Jul 02, 2021
<u>AP</u>		MSN	<u>50MG/VIAL</u>	<u>A212265 001</u>	Aug 28, 2019
<u>AP</u>		NIVAGEN PHARMS INC	<u>50MG/VIAL</u>	<u>A212117 001</u>	Dec 07, 2020
<u>AP</u>		NOVAST LABS	<u>50MG/VIAL</u>	<u>A210984 001</u>	Sep 16, 2019
<u>AP</u>		PHARMASCIENCE INC	<u>50MG/VIAL</u>	<u>A204607 001</u>	May 31, 2017
<u>AP</u>		QILU PHARM HAINAN	<u>50MG/VIAL</u>	<u>A212826 001</u>	Apr 12, 2021
<u>AP</u>		SAGENT PHARMS INC	<u>50MG/VIAL</u>	<u>A207100 001</u>	Mar 16, 2018
<u>AP</u>		SANDOZ	<u>50MG/VIAL</u>	<u>A202969 001</u>	Aug 28, 2014
<u>AP</u>		WOCKHARDT BIO AG	<u>50MG/VIAL</u>	<u>A209056 001</u>	Apr 09, 2019
<u>AP</u>		ZYDUS PHARMS	<u>50MG/VIAL</u>	<u>A214486 001</u>	Nov 19, 2021
<u>POWDER; INTRAVENOUS</u>					
<u>DECITABINE</u>					
	+	SUN PHARM	50MG/VIAL	N205582 001	Jan 28, 2014

DEFERASIROX

GRANULE; ORAL

DEFERASIROX

<u>AB</u>		ALKEM LABS LTD	<u>90MG</u>	<u>A213374 001</u>	Jul 14, 2020
<u>AB</u>			<u>180MG</u>	<u>A213374 002</u>	Jul 14, 2020
<u>AB</u>			<u>360MG</u>	<u>A213374 003</u>	Jul 14, 2020
<u>AB</u>		ANNORA PHARMA	<u>90MG</u>	<u>A216229 003</u>	Mar 16, 2023
<u>AB</u>			<u>180MG</u>	<u>A216229 001</u>	Sep 22, 2022
<u>AB</u>			<u>360MG</u>	<u>A216229 002</u>	Sep 22, 2022
<u>AB</u>		AUCTA	<u>90MG</u>	<u>A214559 001</u>	Mar 09, 2021

PRESCRIPTION DRUG PRODUCT LIST

DEFERASIROX

GRANULE; ORAL

DEFERASIROX

<u>AB</u>		<u>180MG</u>	<u>A214559 002</u>	Mar 09, 2021
<u>AB</u>		<u>360MG</u>	<u>A214559 003</u>	Mar 09, 2021
<u>AB</u>	CIPLA	<u>90MG</u>	<u>A215026 001</u>	Feb 23, 2022
<u>AB</u>		<u>180MG</u>	<u>A215026 002</u>	Feb 23, 2022
<u>AB</u>		<u>360MG</u>	<u>A215026 003</u>	Feb 23, 2022
<u>AB</u>	MSN	<u>90MG</u>	<u>A214650 003</u>	Apr 20, 2022
<u>AB</u>		<u>180MG</u>	<u>A214650 001</u>	Mar 17, 2021
<u>AB</u>		<u>360MG</u>	<u>A214650 002</u>	Mar 17, 2021
<u>AB</u>	TEVA PHARMS USA	<u>90MG</u>	<u>A214180 001</u>	Nov 19, 2021
<u>AB</u>		<u>180MG</u>	<u>A214180 002</u>	Nov 19, 2021
<u>AB</u>		<u>360MG</u>	<u>A214180 003</u>	Nov 19, 2021

JADENU SPRINKLE

<u>AB</u>	+	NOVARTIS	<u>90MG</u>	<u>N207968 001</u>	May 18, 2017
<u>AB</u>	+		<u>180MG</u>	<u>N207968 002</u>	May 18, 2017
<u>AB</u>	+	!	<u>360MG</u>	<u>N207968 003</u>	May 18, 2017

TABLET; ORAL

DEFERASIROX

<u>AB</u>		ACTAVIS ELIZABETH	<u>90MG</u>	<u>A208697 001</u>	Dec 13, 2019
<u>AB</u>			<u>180MG</u>	<u>A208697 002</u>	Dec 13, 2019
<u>AB</u>			<u>360MG</u>	<u>A208697 003</u>	Dec 13, 2019
<u>AB</u>		ALEMBIC	<u>90MG</u>	<u>A211824 001</u>	Nov 20, 2019
<u>AB</u>			<u>180MG</u>	<u>A211824 003</u>	Jun 15, 2020
<u>AB</u>			<u>360MG</u>	<u>A211824 002</u>	Nov 20, 2019
<u>AB</u>		ALKEM LABS LTD	<u>90MG</u>	<u>A210555 001</u>	Mar 30, 2020
<u>AB</u>			<u>180MG</u>	<u>A210555 003</u>	Jul 02, 2020
<u>AB</u>			<u>360MG</u>	<u>A210555 002</u>	Mar 30, 2020
<u>AB</u>		ANNORA PHARMA	<u>90MG</u>	<u>A214341 001</u>	May 14, 2021
<u>AB</u>			<u>180MG</u>	<u>A214341 002</u>	May 14, 2021
<u>AB</u>			<u>360MG</u>	<u>A214341 003</u>	May 14, 2021
<u>AB</u>		AUROBINDO PHARMA LTD	<u>90MG</u>	<u>A214474 001</u>	Oct 16, 2023
<u>AB</u>			<u>180MG</u>	<u>A214474 002</u>	Oct 16, 2023
<u>AB</u>			<u>360MG</u>	<u>A214474 003</u>	Oct 16, 2023
<u>AB</u>		CHARTWELL RX	<u>90MG</u>	<u>A212669 001</u>	May 27, 2021
<u>AB</u>			<u>180MG</u>	<u>A212669 002</u>	May 27, 2021
<u>AB</u>			<u>360MG</u>	<u>A212669 003</u>	May 27, 2021
<u>AB</u>		MSN	<u>90MG</u>	<u>A210945 001</u>	Nov 20, 2019
<u>AB</u>			<u>180MG</u>	<u>A210945 003</u>	Jun 16, 2020
<u>AB</u>			<u>360MG</u>	<u>A210945 002</u>	Nov 20, 2019
<u>AB</u>		NORVIUM BIOSCIENCE	<u>90MG</u>	<u>A211395 001</u>	Mar 22, 2024
<u>AB</u>			<u>180MG</u>	<u>A211395 002</u>	Mar 22, 2024
<u>AB</u>			<u>360MG</u>	<u>A211395 003</u>	Mar 22, 2024
<u>AB</u>		PIRAMAL	<u>90MG</u>	<u>A212995 001</u>	Dec 30, 2019
<u>AB</u>			<u>180MG</u>	<u>A212995 003</u>	Jun 15, 2020
<u>AB</u>			<u>360MG</u>	<u>A212995 002</u>	Dec 30, 2019
<u>AB</u>		SUN PHARM	<u>90MG</u>	<u>A211641 001</u>	Jan 02, 2020
<u>AB</u>			<u>180MG</u>	<u>A211641 003</u>	Jun 15, 2020
<u>AB</u>			<u>360MG</u>	<u>A211641 002</u>	Jan 02, 2020
<u>AB</u>		ZYDUS PHARMS	<u>90MG</u>	<u>A211383 001</u>	Nov 20, 2019
<u>AB</u>			<u>180MG</u>	<u>A211383 003</u>	Jun 15, 2020
<u>AB</u>			<u>360MG</u>	<u>A211383 002</u>	Nov 20, 2019

JADENU

<u>AB</u>	+	NOVARTIS PHARMS CORP	<u>90MG</u>	<u>N206910 001</u>	Mar 30, 2015
<u>AB</u>	+		<u>180MG</u>	<u>N206910 002</u>	Mar 30, 2015
<u>AB</u>	+	!	<u>360MG</u>	<u>N206910 003</u>	Mar 30, 2015

TABLET, FOR SUSPENSION; ORAL

DEFERASIROX

<u>AB</u>		ACTAVIS ELIZABETH	<u>125MG</u>	<u>A203560 001</u>	Jan 26, 2016
<u>AB</u>			<u>250MG</u>	<u>A203560 002</u>	Jan 26, 2016
<u>AB</u>			<u>500MG</u>	<u>A203560 003</u>	Jan 26, 2016
<u>AB</u>		ALEMBIC	<u>125MG</u>	<u>A210060 001</u>	Nov 20, 2019
<u>AB</u>			<u>250MG</u>	<u>A210060 002</u>	Nov 20, 2019
<u>AB</u>			<u>500MG</u>	<u>A210060 003</u>	Nov 20, 2019
<u>AB</u>		ALKEM LABS LTD	<u>125MG</u>	<u>A210519 001</u>	Nov 20, 2019
<u>AB</u>			<u>250MG</u>	<u>A210519 002</u>	Nov 20, 2019
<u>AB</u>			<u>500MG</u>	<u>A210519 003</u>	Nov 20, 2019
<u>AB</u>		BIONPHARMA	<u>125MG</u>	<u>A210920 001</u>	Nov 20, 2019
<u>AB</u>			<u>250MG</u>	<u>A210920 002</u>	Nov 20, 2019
<u>AB</u>			<u>500MG</u>	<u>A210920 003</u>	Nov 20, 2019

PRESCRIPTION DRUG PRODUCT LIST

DEFERASIROX

TABLET, FOR SUSPENSION;ORAL

DEFERASIROX

<u>AB</u>	GLENMARK SPECLT	<u>125MG</u>	<u>A209433 001</u>	Jan 06, 2020
<u>AB</u>		<u>250MG</u>	<u>A209433 002</u>	Jan 06, 2020
<u>AB</u>		<u>500MG</u>	<u>A209433 003</u>	Jan 06, 2020
<u>AB</u>	MSN	<u>125MG</u>	<u>A209878 001</u>	Nov 20, 2019
<u>AB</u>		<u>250MG</u>	<u>A209878 002</u>	Nov 20, 2019
<u>AB</u>		<u>500MG</u>	<u>A209878 003</u>	Nov 20, 2019
<u>AB</u>	SUN PHARM	<u>125MG</u>	<u>A209782 001</u>	Nov 20, 2019
<u>AB</u>		<u>250MG</u>	<u>A209782 002</u>	Nov 20, 2019
<u>AB</u>		<u>500MG</u>	<u>A209782 003</u>	Nov 20, 2019
<u>AB</u>	TEVA PHARMS USA	<u>125MG</u>	<u>A207124 001</u>	Sep 23, 2022
<u>AB</u>		<u>250MG</u>	<u>A207124 002</u>	Sep 23, 2022
<u>AB</u>		<u>500MG</u>	<u>A207124 003</u>	Sep 23, 2022
<u>AB</u>	TORRENT	<u>125MG</u>	<u>A209426 001</u>	Nov 05, 2024
<u>AB</u>		<u>250MG</u>	<u>A209426 002</u>	Nov 05, 2024
<u>AB</u>		<u>500MG</u>	<u>A209426 003</u>	Nov 05, 2024
<u>EXJADE</u>				
<u>AB</u>	+ NOVARTIS	<u>125MG</u>	<u>N021882 001</u>	Nov 02, 2005
<u>AB</u>	+	<u>250MG</u>	<u>N021882 002</u>	Nov 02, 2005
<u>AB</u>	+!	<u>500MG</u>	<u>N021882 003</u>	Nov 02, 2005

DEFERIPRONE

SOLUTION;ORAL

FERRIPROX

+! CHIESI

100MG/ML

N208030 001 Sep 09, 2015

TABLET;ORAL

DEFERIPRONE

<u>AB</u>	HIKMA	<u>500MG</u>	<u>A213239 001</u>	Mar 29, 2021
<u>AB</u>	!	<u>1GM</u>	<u>A213239 002</u>	Feb 08, 2022
<u>AB</u>	TARO	<u>500MG</u>	<u>A208800 001</u>	Feb 08, 2019
<u>AB</u>		<u>1GM</u>	<u>A208800 002</u>	Nov 22, 2023

FERRIPROX

<u>AB</u>	+ CHIESI	<u>500MG</u>	<u>N021825 001</u>	Oct 14, 2011
<u>AB</u>	+	<u>1GM</u>	<u>N021825 002</u>	Jul 25, 2019
	+!	1GM	N212269 001	May 19, 2020

DEFEROXAMINE MESYLATE

INJECTABLE; INJECTION

DEFEROXAMINE MESYLATE

<u>AP</u>	FRESENIUS KABI USA	<u>500MG/VIAL</u>	<u>A078718 001</u>	Sep 15, 2009
<u>AP</u>		<u>2GM/VIAL</u>	<u>A078718 002</u>	Sep 15, 2009
<u>AP</u>	GLAND PHARMA LTD	<u>500MG/VIAL</u>	<u>A207384 001</u>	Sep 29, 2017
<u>AP</u>		<u>2GM/VIAL</u>	<u>A207384 002</u>	Sep 29, 2017
<u>AP</u>	HIKMA	<u>500MG/VIAL</u>	<u>A078086 001</u>	May 30, 2007
<u>AP</u>		<u>2GM/VIAL</u>	<u>A078086 002</u>	May 30, 2007
<u>AP</u>	HOSPIRA	<u>500MG/VIAL</u>	<u>A076019 001</u>	Mar 17, 2004
<u>AP</u>	!	<u>2GM/VIAL</u>	<u>A076019 002</u>	Mar 17, 2004
<u>DESFERAL</u>				
<u>AP</u>	+! NOVARTIS	<u>500MG/VIAL</u>	<u>N016267 001</u>	

DEFIBROTIDE SODIUM

SOLUTION; INTRAVENOUS

DEFITELIO

+! JAZZ PHARMS INC

200MG/2.5ML (80MG/ML)

N208114 001 Mar 30, 2016

DEFLAZACORT

SUSPENSION;ORAL

DEFLAZACORT

<u>AB</u>	TRIS PHARMA INC	<u>22.75MG/ML</u>	<u>A217813 001</u>	Apr 25, 2024
<u>EMFLAZA</u>				
<u>AB</u>	+! PTC THERAP	<u>22.75MG/ML</u>	<u>N208685 001</u>	Feb 09, 2017

TABLET;ORAL

DEFLAZACORT

<u>AB</u>	AUROBINDO PHARMA LTD	<u>6MG</u>	<u>A217123 001</u>	Feb 09, 2024
<u>AB</u>		<u>18MG</u>	<u>A217123 002</u>	Feb 09, 2024
<u>AB</u>		<u>30MG</u>	<u>A217123 003</u>	Feb 09, 2024
<u>AB</u>		<u>36MG</u>	<u>A217123 004</u>	Feb 09, 2024
<u>AB</u>	UPSHER SMITH LABS	<u>6MG</u>	<u>A216720 001</u>	Nov 05, 2024
<u>AB</u>		<u>18MG</u>	<u>A216720 002</u>	Nov 05, 2024
<u>AB</u>		<u>30MG</u>	<u>A216720 003</u>	Nov 05, 2024
<u>AB</u>		<u>36MG</u>	<u>A216720 004</u>	Nov 05, 2024

PRESCRIPTION DRUG PRODUCT LIST

DEFLAZACORT

TABLET; ORAL

EMFLAZA

AB	+	PTC THERAP	6MG	N208684 001	Feb 09, 2017
AB	+		18MG	N208684 002	Feb 09, 2017
AB	+		30MG	N208684 003	Feb 09, 2017
AB	+		36MG	N208684 004	Feb 09, 2017

DEGARELIX ACETATE

POWDER; SUBCUTANEOUS

FIRMAGON

+	FERRING	EQ 80MG BASE/VIAL	N022201 001	Dec 24, 2008
+	!	EQ 120MG BASE/VIAL	N022201 002	Dec 24, 2008

DELAFLOXACIN MEGLUMINE

POWDER; INTRAVENOUS

BAXDELA

+	!	MELINTA	EQ 300MG BASE/VIAL	N208611 001	Jun 19, 2017
---	---	---------	--------------------	-------------	--------------

TABLET; ORAL

BAXDELA

+	!	MELINTA	EQ 450MG BASE	N208610 001	Jun 19, 2017
---	---	---------	---------------	-------------	--------------

DEMECLOCYCLINE HYDROCHLORIDE

TABLET; ORAL

DEMECLOCYCLINE HYDROCHLORIDE

AB		AMNEAL PHARM	150MG	A065425 001	Feb 27, 2008
AB	!		300MG	A065425 002	Feb 27, 2008
AB		EPIC PHARMA LLC	150MG	A065389 001	Dec 01, 2008
AB			150MG	A065447 001	Aug 18, 2015
AB			300MG	A065389 002	Dec 01, 2008
AB			300MG	A065447 002	Aug 18, 2015

DEOXYCHOLIC ACID

SOLUTION; SUBCUTANEOUS

DEOXYCHOLIC ACID

AP		WILSHIRE PHARMS INC	20MG/2ML (10MG/ML)	A212296 001	Apr 02, 2021
-----------	--	---------------------	---------------------------	--------------------	--------------

KYBELLA

AP	+	!	ABEVIE	20MG/2ML (10MG/ML)	N206333 001	Apr 29, 2015
-----------	---	---	--------	---------------------------	--------------------	--------------

DESFLURANE

LIQUID; INHALATION

DESFLURANE

AN		SHANGHAI HENGRUI	100%	A208234 001	Feb 26, 2018
-----------	--	------------------	-------------	--------------------	--------------

SUPRANE

AN	+	!	BAXTER HLTHCARE	100%	N020118 001	Sep 18, 1992
-----------	---	---	-----------------	-------------	--------------------	--------------

DESIPRAMINE HYDROCHLORIDE

TABLET; ORAL

DESIPRAMINE HYDROCHLORIDE

AB		ACTAVIS TOTOWA	10MG	A074430 001	Feb 09, 1996
AB			25MG	A071601 001	Jun 05, 1987
AB			50MG	A071588 001	Jun 05, 1987
AB			75MG	A071602 001	Oct 05, 1987
AB			100MG	A071766 001	Oct 05, 1987
AB			150MG	A074430 002	Feb 09, 1996
AB		ALEMBIC	10MG	A209785 001	Jul 07, 2021
AB			25MG	A209785 002	Jul 07, 2021
AB			50MG	A209785 003	Jul 07, 2021
AB			75MG	A209785 004	Jul 07, 2021
AB			100MG	A209785 005	Jul 07, 2021
AB			150MG	A209785 006	Jul 07, 2021
AB		AMNEAL PHARMS CO	10MG	A208105 001	Mar 17, 2016
AB			25MG	A208105 002	Mar 17, 2016
AB			50MG	A208105 003	Mar 17, 2016
AB			75MG	A208105 004	Mar 17, 2016
AB			100MG	A208105 005	Mar 17, 2016
AB			150MG	A208105 006	Mar 17, 2016
AB		CHARTWELL RX	10MG	A072103 002	May 24, 1988
AB			25MG	A072103 003	May 24, 1988
AB			50MG	A072103 004	May 24, 1988
AB			75MG	A072103 005	Jun 20, 1988
AB			100MG	A072103 001	Jun 20, 1988
AB			150MG	A072103 006	Jun 20, 1988
AB		HERITAGE	10MG	A207433 001	May 05, 2016
AB			25MG	A207433 002	May 05, 2016

PRESCRIPTION DRUG PRODUCT LIST

DESIPRAMINE HYDROCHLORIDE

TABLET; ORAL

DESIPRAMINE HYDROCHLORIDE

<u>AB</u>		<u>50MG</u>	<u>A207433</u>	<u>003</u>	May 05, 2016
<u>AB</u>		<u>75MG</u>	<u>A207433</u>	<u>004</u>	May 05, 2016
<u>AB</u>		<u>100MG</u>	<u>A207433</u>	<u>005</u>	May 05, 2016
<u>AB</u>		<u>150MG</u>	<u>A207433</u>	<u>006</u>	May 05, 2016
<u>AB</u>	NOVAST LABS	<u>10MG</u>	<u>A204963</u>	<u>001</u>	Dec 26, 2017
<u>AB</u>		<u>25MG</u>	<u>A204963</u>	<u>002</u>	Dec 26, 2017
<u>AB</u>		<u>50MG</u>	<u>A204963</u>	<u>003</u>	Dec 26, 2017
<u>AB</u>		<u>75MG</u>	<u>A204963</u>	<u>004</u>	Dec 26, 2017
<u>AB</u>		<u>100MG</u>	<u>A204963</u>	<u>005</u>	Dec 26, 2017
<u>AB</u>		<u>150MG</u>	<u>A204963</u>	<u>006</u>	Dec 26, 2017

NORPRAMIN

<u>AB</u>	+	VALIDUS PHARMS	<u>10MG</u>	<u>N014399</u>	<u>007</u>	Feb 11, 1982
<u>AB</u>	+		<u>25MG</u>	<u>N014399</u>	<u>001</u>	
<u>AB</u>	+		<u>50MG</u>	<u>N014399</u>	<u>003</u>	
<u>AB</u>	+		<u>75MG</u>	<u>N014399</u>	<u>004</u>	
<u>AB</u>	+		<u>100MG</u>	<u>N014399</u>	<u>005</u>	
<u>AB</u>	+		<u>150MG</u>	<u>N014399</u>	<u>006</u>	

DES Loratadine

TABLET; ORAL

CLARINEX

<u>AB</u>	+	ORGANON	<u>5MG</u>	<u>N021165</u>	<u>001</u>	Dec 21, 2001
<u>DES Loratadine</u>						
<u>AB</u>		BELCHER PHARMS	<u>5MG</u>	<u>A078355</u>	<u>001</u>	Apr 19, 2012
<u>AB</u>		DR REDDYS LABS LTD	<u>5MG</u>	<u>A078365</u>	<u>001</u>	Mar 08, 2011
<u>AB</u>		LUPIN PHARMS	<u>5MG</u>	<u>A078352</u>	<u>001</u>	Oct 25, 2010
<u>AB</u>		ORBION PHARMS	<u>5MG</u>	<u>A078357</u>	<u>001</u>	Feb 19, 2010

TABLET, ORALLY DISINTEGRATING; ORAL

DES Loratadine

		REDDYS	2.5MG	A078367	001	Jul 12, 2010
!			5MG	A078367	002	Jul 12, 2010

DES Loratadine; Pseudoephedrine Sulfate

TABLET, EXTENDED RELEASE; ORAL

CLARINEX-D 12 HOUR

+	!	ORGANON LLC	2.5MG;120MG	N021313	001	Feb 01, 2006
---	---	-------------	-------------	---------	-----	--------------

DES Loratadine AND Pseudoephedrine Sulfate 24 HOUR

!		DR REDDYS LABS LTD	5MG;240MG	A078366	001	Apr 26, 2011
---	--	--------------------	-----------	---------	-----	--------------

Desmopressin Acetate

INJECTABLE; INJECTION

DDAVP

<u>AP</u>	+	FERRING PHARMS INC	<u>0.004MG/ML</u>	<u>N018938</u>	<u>001</u>	Mar 30, 1984
-----------	---	--------------------	-------------------	----------------	------------	--------------

Desmopressin Acetate

<u>AP</u>		DR REDDYS	<u>0.004MG/ML</u>	<u>A215961</u>	<u>001</u>	Aug 19, 2022
<u>AP</u>		GLAND	<u>0.004MG/ML</u>	<u>A216904</u>	<u>001</u>	Mar 20, 2023
<u>AP</u>		GLAND PHARMA LTD	<u>0.004MG/ML</u>	<u>A216922</u>	<u>001</u>	Nov 16, 2022
<u>AP</u>		MEITHEAL	<u>0.004MG/ML</u>	<u>A074888</u>	<u>001</u>	Oct 15, 1997
<u>AP</u>		SAGENT PHARMS INC	<u>0.004MG/ML</u>	<u>A204695</u>	<u>001</u>	Aug 22, 2017
<u>AP</u>			<u>0.004MG/ML</u>	<u>A204751</u>	<u>001</u>	Aug 22, 2017
<u>AP</u>		UBI	<u>0.004MG/ML</u>	<u>A210218</u>	<u>001</u>	Feb 14, 2020
<u>AP</u>			<u>0.004MG/ML</u>	<u>A210223</u>	<u>001</u>	Sep 17, 2020

SPRAY, METERED; NASAL

Desmopressin Acetate

<u>AB</u>	!	BAUSCH	<u>0.01MG/SPRAY</u>	<u>A074830</u>	<u>001</u>	Jan 25, 1999
-----------	---	--------	---------------------	----------------	------------	--------------

Desmopressin Acetate (Needs No Refrigeration)

<u>AB</u>	!	APOTEX	<u>0.01MG/SPRAY</u>	<u>A076703</u>	<u>001</u>	Jan 27, 2005
<u>AB</u>		ZYDUS PHARMS	<u>0.01MG/SPRAY</u>	<u>A091345</u>	<u>001</u>	Oct 03, 2017

TABLET; ORAL

DDAVP

<u>AB</u>	+	FERRING PHARMS INC	<u>0.1MG</u>	<u>N019955</u>	<u>001</u>	Sep 06, 1995
<u>AB</u>	+		<u>0.2MG</u>	<u>N019955</u>	<u>002</u>	Sep 06, 1995

Desmopressin Acetate

<u>AB</u>		ABHAI LLC	<u>0.1MG</u>	<u>A210371</u>	<u>001</u>	Jan 28, 2019
<u>AB</u>			<u>0.2MG</u>	<u>A210371</u>	<u>002</u>	Jan 28, 2019
<u>AB</u>		APOTEX INC	<u>0.1MG</u>	<u>A077414</u>	<u>001</u>	Mar 07, 2006
<u>AB</u>			<u>0.2MG</u>	<u>A077414</u>	<u>002</u>	Mar 07, 2006
<u>AB</u>		AUROBINDO PHARMA	<u>0.1MG</u>	<u>A213095</u>	<u>001</u>	Mar 21, 2024
<u>AB</u>			<u>0.2MG</u>	<u>A213095</u>	<u>002</u>	Mar 21, 2024
<u>AB</u>		GLENMARK PHARMS LTD	<u>0.1MG</u>	<u>A201831</u>	<u>001</u>	May 28, 2015
<u>AB</u>			<u>0.2MG</u>	<u>A201831</u>	<u>002</u>	May 28, 2015

PRESCRIPTION DRUG PRODUCT LIST

DESMOPRESSIN ACETATE

TABLET;ORAL

DESMOPRESSIN ACETATE

<u>AB</u>	HERITAGE PHARMA	<u>0.1MG</u>	<u>A207880 001</u>	May 26, 2017
<u>AB</u>		<u>0.2MG</u>	<u>A207880 002</u>	May 26, 2017
<u>AB</u>	NOVAST LABS	<u>0.1MG</u>	<u>A208357 001</u>	Jun 06, 2019
<u>AB</u>		<u>0.2MG</u>	<u>A208357 002</u>	Jun 06, 2019

DESOGESTREL; ETHINYL ESTRADIOL

TABLET;ORAL-28

BEKYREE

<u>AB</u>	LUPIN LTD	<u>0.15MG,N/A;0.02MG,0.01MG</u>	<u>A202226 001</u>	Aug 12, 2015
-----------	-----------	---------------------------------	--------------------	--------------

CYCLESSA

<u>AB</u>	+! ASPEN GLOBAL INC	<u>0.1MG,0.125MG,0.15MG;0.025MG,0.025MG,0.025MG</u>	<u>N021090 001</u>	Dec 20, 2000
-----------	---------------------	---	--------------------	--------------

DESOGESTREL AND ETHINYL ESTRADIOL

<u>AB</u>	! DURAMED PHARMS BARR	<u>0.15MG;0.03MG</u>	<u>A075256 002</u>	Aug 12, 1999
<u>AB</u>	NAARI PTE LTD	<u>0.15MG,N/A;0.02MG,0.01MG</u>	<u>A209170 001</u>	Jun 05, 2017
<u>AB</u>		<u>0.15MG;0.03MG</u>	<u>A207067 001</u>	Sep 13, 2018
<u>AB</u>	NOVAST LABS	<u>0.15MG;0.03MG</u>	<u>A091234 001</u>	Jul 12, 2013
<u>AB</u>	WATSON LABS	<u>0.15MG;0.03MG</u>	<u>A076915 001</u>	Jul 29, 2005
<u>AB</u>	XIROMED	<u>0.15MG,N/A;0.02MG,0.01MG</u>	<u>A202296 001</u>	Aug 30, 2013

ENSKYCE

<u>AB</u>	LUPIN LTD	<u>0.15MG;0.03MG</u>	<u>A201887 001</u>	Mar 07, 2013
-----------	-----------	----------------------	--------------------	--------------

ISIBLOOM

<u>AB</u>	XIROMED	<u>0.15MG;0.03MG</u>	<u>A202789 001</u>	Aug 12, 2015
-----------	---------	----------------------	--------------------	--------------

KALLIGA

<u>AB</u>	AUROBINDO PHARMA	<u>0.15MG;0.03MG</u>	<u>A207081 001</u>	May 17, 2017
-----------	------------------	----------------------	--------------------	--------------

KARIVA

<u>AB</u>	! BARR	<u>0.15MG,N/A;0.02MG,0.01MG</u>	<u>A075863 001</u>	Apr 05, 2002
-----------	--------	---------------------------------	--------------------	--------------

PIMTREA

<u>AB</u>	NOVAST LABS	<u>0.15MG,N/A;0.02MG,0.01MG</u>	<u>A091247 001</u>	Aug 01, 2013
-----------	-------------	---------------------------------	--------------------	--------------

SIMLIYA

<u>AB</u>	AUROBINDO PHARMA	<u>0.15MG,N/A;0.02MG,0.01MG</u>	<u>A206853 001</u>	Mar 22, 2017
-----------	------------------	---------------------------------	--------------------	--------------

VELIVET

<u>AB</u>	DURAMED PHARMS BARR	<u>0.1MG,0.125MG,0.15MG;0.025MG,0.025MG,0.025MG</u>	<u>A076455 001</u>	Feb 24, 2004
-----------	---------------------	---	--------------------	--------------

VIORELE

<u>AB</u>	GLENMARK PHARMS LTD	<u>0.15MG,N/A;0.02MG,0.01MG</u>	<u>A091346 001</u>	Apr 02, 2012
-----------	---------------------	---------------------------------	--------------------	--------------

VOLNEA

<u>AB</u>	XIROMED	<u>0.15MG,N/A;0.02MG,0.01MG</u>	<u>A202689 001</u>	Sep 09, 2016
-----------	---------	---------------------------------	--------------------	--------------

DESONIDE

CREAM;TOPICAL

DESONIDE

<u>AB</u>	ALEMBIC	<u>0.05%</u>	<u>A214396 001</u>	Dec 08, 2022
<u>AB</u>	CADILA	<u>0.05%</u>	<u>A210198 001</u>	Nov 20, 2019
<u>AB</u>	COSETTE	<u>0.05%</u>	<u>A074027 001</u>	Sep 28, 1992
<u>AB</u>	GLENMARK SPECLT	<u>0.05%</u>	<u>A209729 001</u>	Jul 24, 2017
<u>AB</u>	+! PADAGIS US	<u>0.05%</u>	<u>N017010 001</u>	
<u>AB</u>	TARO	<u>0.05%</u>	<u>A073548 001</u>	Jun 30, 1992

DESOWEN

<u>AB</u>	GALDERMA LABS LP	<u>0.05%</u>	<u>N019048 001</u>	Dec 14, 1984
-----------	------------------	--------------	--------------------	--------------

GEL;TOPICAL

DESONIDE

! CINTEX SVCS

0.05% A202470 001 May 11, 2020

LOTION;TOPICAL

DESONIDE

<u>AB</u>	ALEMBIC	<u>0.05%</u>	<u>A213632 001</u>	Aug 24, 2020
<u>AB</u>	FOUGERA PHARMS	<u>0.05%</u>	<u>A075860 001</u>	Mar 19, 2002
<u>AB</u>	GLENMARK SPECLT	<u>0.05%</u>	<u>A209494 001</u>	Sep 26, 2017
<u>AB</u>	TARO	<u>0.05%</u>	<u>A202161 001</u>	Oct 31, 2014

DESOWEN

<u>AB</u>	! GALDERMA LABS LP	<u>0.05%</u>	<u>A072354 001</u>	Jan 24, 1992
-----------	--------------------	--------------	--------------------	--------------

OINTMENT;TOPICAL

DESONIDE

<u>AB</u>	ALEMBIC	<u>0.05%</u>	<u>A212473 001</u>	Oct 23, 2019
<u>AB</u>	ENCUBE ETHICALS	<u>0.05%</u>	<u>A210998 001</u>	Jan 30, 2019
<u>AB</u>	FOUGERA PHARMS	<u>0.05%</u>	<u>A075751 001</u>	Mar 12, 2001
<u>AB</u>	GLENMARK PHARMS LTD	<u>0.05%</u>	<u>A209996 001</u>	Sep 15, 2017
<u>AB</u>	+! PADAGIS US	<u>0.05%</u>	<u>N017426 001</u>	
<u>AB</u>	TARO	<u>0.05%</u>	<u>A074254 001</u>	Aug 03, 1994

PRESCRIPTION DRUG PRODUCT LIST

DESONIDE

OINTMENT; TOPICAL

DESOWEN

AB GALDERMA LABS LP **0.05%** **A071425 001** Jun 15, 1988

DESOXIMETASONE

CREAM; TOPICAL

DESOXIMETASONE

AB ACTAVIS MID **0.25%** **A205082 001** Sep 04, 2015

ATLANTIC

AB FOUGERA PHARMS **0.25%** **A078369 001** Jun 29, 2010

AB LUPIN **0.05%** **A208163 001** Jan 10, 2017

AB **0.25%** **A208164 001** Jan 09, 2017

AB PADAGIS ISRAEL **0.25%** **A076510 001** Jul 01, 2003

AB RISING **0.05%** **A210980 001** Dec 21, 2018

AB **0.25%** **A205594 001** Jul 02, 2018

AB ZYDUS LIFESCIENCES **0.25%** **A205620 001** Sep 28, 2018

TOPICORT

AB ! TARO **0.05%** **A073210 001** Nov 30, 1990

AB ! **0.25%** **A073193 001** Nov 30, 1990

GEL; TOPICAL

DESOXIMETASONE

AB EPIC PHARMA LLC **0.05%** **A090727 001** Mar 10, 2011

AB PADAGIS US **0.05%** **A077552 001** Jan 09, 2006

AB RISING **0.05%** **A204675 001** Aug 12, 2016

TOPICORT

AB ! TARO **0.05%** **A074904 001** Jul 14, 1998

OINTMENT; TOPICAL

DESOXIMETASONE

AB ACTAVIS MID **0.25%** **A204965 001** Nov 07, 2016

ATLANTIC

AB EPIC PHARMA LLC **0.25%** **A201005 001** Apr 24, 2014

AB FOUGERA PHARMS **0.25%** **A078657 001** Sep 28, 2012

AB ! GLENMARK PHARMS LTD **0.25%** **A202838 001** Sep 20, 2013

AB LUPIN **0.05%** **A208044 001** Dec 12, 2016

AB **0.25%** **A208104 001** Dec 01, 2016

AB NOVEL LABS INC **0.25%** **A206792 001** May 10, 2016

AB PADAGIS ISRAEL **0.25%** **A077770 001** Apr 20, 2015

AB RISING **0.25%** **A204272 001** Nov 30, 2016

AB THE J MOLNER **0.05%** **A209973 001** Oct 23, 2018

AB ZYDUS LIFESCIENCES **0.25%** **A205206 001** Sep 19, 2017

TOPICORT

AB +! TARO **0.05%** **N018594 001** Jan 17, 1985

SPRAY; TOPICAL

DESOXIMETASONE

AT LUPIN **0.25%** **A208124 001** Mar 16, 2018

AT PADAGIS ISRAEL **0.25%** **A206441 001** Jan 20, 2017

TOPICORT

AT +! TARO **0.25%** **N204141 001** Apr 11, 2013

DESVENLAFAXINE

TABLET, EXTENDED RELEASE; ORAL

DESVENLAFAXINE

+ ALEMBIC PHARMS LTD 50MG N204150 001 Mar 04, 2013

+! 100MG N204150 002 Mar 04, 2013

DESVENLAFAXINE SUCCINATE

TABLET, EXTENDED RELEASE; ORAL

DESVENLAFAXINE SUCCINATE

AB ACTAVIS LABS FL **EQ 25MG BASE** **A204065 001** Jul 29, 2016

AB **EQ 50MG BASE** **A204065 002** Jul 29, 2016

AB **EQ 100MG BASE** **A204065 003** Jul 29, 2016

AB ALEMBIC **EQ 25MG BASE** **A204003 003** Sep 14, 2018

AB **EQ 50MG BASE** **A204003 001** Jun 29, 2015

AB **EQ 100MG BASE** **A204003 002** Jun 29, 2015

AB HIKMA **EQ 25MG BASE** **A204082 002** Aug 28, 2017

AB **EQ 50MG BASE** **A204082 001** Feb 16, 2016

AB **EQ 100MG BASE** **A204083 001** Feb 16, 2016

AB INTELLIPHARMACEUTIC **EQ 50MG BASE** **A204805 001** May 07, 2019

S

AB **EQ 100MG BASE** **A204805 002** May 07, 2019

AB LUPIN LTD **EQ 25MG BASE** **A204172 003** Apr 13, 2022

AB **EQ 50MG BASE** **A204172 001** Jun 29, 2015

AB **EQ 100MG BASE** **A204172 002** Jun 29, 2015

PRESCRIPTION DRUG PRODUCT LIST

DESVENLAFAXINE SUCCINATE

TABLET, EXTENDED RELEASE;ORAL

DESVENLAFAXINE SUCCINATE

<u>AB</u>	RUBICON	<u>EQ 25MG BASE</u>	<u>A204028 003</u>	Dec 07, 2023
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A204028 001</u>	Jun 29, 2015
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A204028 002</u>	Jun 29, 2015
<u>AB</u>	YICHANG HUMANWELL	<u>EQ 25MG BASE</u>	<u>A210014 003</u>	Oct 13, 2020
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A210014 001</u>	Oct 01, 2018
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A210014 002</u>	Oct 01, 2018
<u>AB</u>	ZYDUS PHARMS	<u>EQ 25MG BASE</u>	<u>A204020 003</u>	Nov 30, 2022
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A204020 001</u>	Oct 11, 2017
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A204020 002</u>	Oct 11, 2017
<u>PRISTIO</u>				
<u>AB</u>	+ PF PRISM CV	<u>EQ 25MG BASE</u>	<u>N021992 003</u>	Aug 20, 2014
<u>AB</u>	+	<u>EQ 50MG BASE</u>	<u>N021992 001</u>	Feb 29, 2008
<u>AB</u>	+!	<u>EQ 100MG BASE</u>	<u>N021992 002</u>	Feb 29, 2008

DEUCRAVACITINIB

TABLET;ORAL

SOTYKTU

+! BRISTOL

6MG

N214958 001 Sep 09, 2022

DEURUXOLITINIB PHOSPHATE

TABLET;ORAL

LEQSELVI

+! SUN PHARM INDS INC

EQ 8MG BASE

N217900 001 Jul 25, 2024

DEUTETRABENAZINE

TABLET;ORAL

AUSTEDO

<u>AB</u>	+ TEVA BRANDED PHARM	<u>6MG</u>	<u>N208082 001</u>	Apr 03, 2017
<u>AB</u>	+	<u>9MG</u>	<u>N208082 002</u>	Apr 03, 2017
<u>AB</u>	+!	<u>12MG</u>	<u>N208082 003</u>	Apr 03, 2017

DEUTETRABENAZINE

<u>AB</u>	AUROBINDO PHARMA LTD	<u>6MG</u>	<u>A215971 001</u>	Dec 26, 2024
<u>AB</u>		<u>9MG</u>	<u>A215971 002</u>	Dec 26, 2024
<u>AB</u>		<u>12MG</u>	<u>A215971 003</u>	Dec 26, 2024

TABLET, EXTENDED RELEASE;ORAL

AUSTEDO XR

+ TEVA

6MG

N216354 001 Feb 17, 2023

+

12MG

N216354 002 Feb 17, 2023

+

18MG

N216354 008 Jul 01, 2024

+!

24MG

N216354 003 Feb 17, 2023

+

30MG

N216354 004 May 29, 2024

+

36MG

N216354 005 May 29, 2024

+

42MG

N216354 006 May 29, 2024

+

48MG

N216354 007 May 29, 2024

DEUTIVACAFTOR; TEZACAFTOR; VANZACAFTOR CALCIUM

TABLET;ORAL

ALYFTREK

+ VERTEX PHARMS INC

50MG;20MG;EQ 4MG BASE

N218730 001 Dec 20, 2024

+!

125MG;50MG;EQ 10MG BASE

N218730 002 Dec 20, 2024

DEXAMETHASONE

CONCENTRATE;ORAL

DEXAMETHASONE INTENSOL

! HIKMA

1MG/ML

A088252 001 Sep 01, 1983

ELIXIR;ORAL

DEXAMETHASONE

<u>AA</u>	CHARTWELL MOLECULAR	<u>0.5MG/5ML</u>	<u>A091188 001</u>	May 11, 2011
<u>AA</u>	! KANCHAN HLTHCARE	<u>0.5MG/5ML</u>	<u>A084754 001</u>	
<u>AA</u>	RISING	<u>0.5MG/5ML</u>	<u>A090891 001</u>	Jul 12, 2011
IMPLANT;INTRAVITREAL				
OZURDEX				
	+! ABBVIE	0.7MG	N022315 001	Jun 17, 2009
INSERT;OPHTHALMIC				
DEXTENZA				
	+! OCULAR THERAPEUTIX	0.4MG	N208742 001	Nov 30, 2018
SOLUTION;ORAL				
DEXAMETHASONE				
	+! HIKMA	0.5MG/5ML	A088248 001	Sep 01, 1983

PRESCRIPTION DRUG PRODUCT LIST

DEXAMETHASONE

SUSPENSION; INTRAOCULAR				
DEXYCU KIT				
+	!	EYEPOINT PHARMS	9%	N208912 001 Feb 09, 2018
SUSPENSION/DROPS; OPHTHALMIC				
MAXIDEX				
+	!	HARROW EYE	0.1%	N013422 001

TABLET; ORAL

<u>DEXAMETHASONE</u>				
<u>AB</u>		ALVOGEN	<u>4MG</u>	<u>A088481 004</u> Apr 28, 1983
<u>AB</u>			<u>6MG</u>	<u>A088481 001</u> Nov 28, 1983
<u>AB</u>		AMNEAL	<u>2MG</u>	<u>A216295 001</u> Sep 08, 2022
<u>AB</u>			<u>4MG</u>	<u>A215106 001</u> Oct 14, 2021
<u>AB</u>			<u>6MG</u>	<u>A215106 002</u> Oct 14, 2021
<u>AB</u>		APOTEX	<u>0.5MG</u>	<u>A217695 001</u> Aug 23, 2023
<u>AB</u>			<u>0.75MG</u>	<u>A217695 002</u> Aug 23, 2023
<u>AB</u>			<u>1MG</u>	<u>A217695 003</u> Aug 23, 2023
<u>AB</u>			<u>1.5MG</u>	<u>A217695 004</u> Aug 23, 2023
<u>AB</u>			<u>2MG</u>	<u>A217695 005</u> Aug 23, 2023
<u>AB</u>			<u>4MG</u>	<u>A217695 006</u> Aug 23, 2023
<u>AB</u>			<u>6MG</u>	<u>A217695 007</u> Aug 23, 2023
<u>AB</u>		BIONPHARMA	<u>2MG</u>	<u>A217538 001</u> Apr 26, 2023
<u>AB</u>			<u>4MG</u>	<u>A217001 001</u> Apr 19, 2023
<u>AB</u>			<u>6MG</u>	<u>A217001 002</u> Apr 19, 2023
<u>AB</u>		COREPHARMA	<u>0.5MG</u>	<u>A218372 001</u> Sep 17, 2024
<u>AB</u>			<u>0.75MG</u>	<u>A218372 002</u> Sep 17, 2024
<u>AB</u>			<u>1MG</u>	<u>A218372 003</u> Sep 17, 2024
<u>AB</u>			<u>1.5MG</u>	<u>A218372 004</u> Sep 17, 2024
<u>AB</u>			<u>2MG</u>	<u>A218372 005</u> Sep 17, 2024
<u>AB</u>			<u>4MG</u>	<u>A218372 006</u> Sep 17, 2024
<u>AB</u>			<u>6MG</u>	<u>A218372 007</u> Sep 17, 2024
<u>AB</u>	+	HIKMA	<u>0.5MG</u>	<u>A084611 001</u>
<u>AB</u>	+		<u>0.75MG</u>	<u>A084613 001</u>
<u>AB</u>	+	!	<u>1MG</u>	<u>A088306 001</u> Sep 15, 1983
<u>AB</u>	+		<u>1.5MG</u>	<u>A084610 001</u>
<u>AB</u>	+	!	<u>2MG</u>	<u>A087916 001</u> Aug 26, 1982
<u>AB</u>	+		<u>4MG</u>	<u>A084612 001</u>
<u>AB</u>	+	!	<u>6MG</u>	<u>A088316 001</u> Sep 15, 1983
<u>AB</u>		LARKEN LABS INC	<u>1.5MG</u>	<u>A201270 001</u> Jul 17, 2017
<u>AB</u>		NOVITIUM PHARMA	<u>0.5MG</u>	<u>A215604 004</u> Oct 05, 2023
<u>AB</u>			<u>0.75MG</u>	<u>A215604 005</u> Oct 05, 2023
<u>AB</u>			<u>1.5MG</u>	<u>A215604 001</u> Aug 08, 2022
<u>AB</u>			<u>2MG</u>	<u>A217696 001</u> May 09, 2023
<u>AB</u>			<u>4MG</u>	<u>A215604 002</u> Aug 08, 2022
<u>AB</u>			<u>6MG</u>	<u>A215604 003</u> Aug 08, 2022
<u>AB</u>		PRASCO	<u>4MG</u>	<u>A080399 002</u> Apr 20, 2022
<u>AB</u>		ZYDUS LIFESCIENCES	<u>0.5MG</u>	<u>A216282 001</u> Feb 07, 2024
<u>AB</u>			<u>0.75MG</u>	<u>A216282 002</u> Feb 07, 2024
<u>AB</u>			<u>1MG</u>	<u>A216284 001</u> May 09, 2024
<u>AB</u>			<u>1.5MG</u>	<u>A216282 003</u> Feb 07, 2024
<u>AB</u>			<u>2MG</u>	<u>A216283 001</u> Feb 07, 2024
<u>AB</u>			<u>4MG</u>	<u>A216282 004</u> Feb 07, 2024
<u>AB</u>			<u>6MG</u>	<u>A216282 005</u> Feb 07, 2024
BP		ALVOGEN	0.5MG	A088481 002 Apr 28, 1983
BP			0.75MG	A088481 003 Apr 28, 1983
HEMADY				
+	!	DEXCEL	20MG	N211379 001 Oct 03, 2019

DEXAMETHASONE SODIUM PHOSPHATE

INJECTABLE; INJECTION				
<u>DEXAMETHASONE SODIUM PHOSPHATE</u>				
<u>AP</u>		AMNEAL	<u>EQ 4MG PHOSPHATE/ML</u>	<u>A208689 001</u> Aug 22, 2018
<u>AP</u>		EUGIA PHARMA	<u>EQ 4MG PHOSPHATE/ML</u>	<u>A206781 001</u> Dec 01, 2015
<u>AP</u>			<u>EQ 10MG PHOSPHATE/ML</u>	<u>A210966 001</u> Jun 05, 2020
<u>AP</u>			<u>EQ 10MG PHOSPHATE/ML</u>	<u>A210967 001</u> Jun 07, 2019
<u>AP</u>	+	FRESENIUS KABI USA	<u>EQ 4MG PHOSPHATE/ML</u>	<u>A084916 001</u>
<u>AP</u>			<u>EQ 4MG PHOSPHATE/ML</u>	<u>A203129 001</u> Sep 30, 2015
<u>AP</u>	!		<u>EQ 10MG PHOSPHATE/ML</u>	<u>A040572 001</u> Apr 22, 2005
<u>AP</u>			<u>EQ 10MG PHOSPHATE/ML</u>	<u>A209192 001</u> Jul 06, 2018
<u>AP</u>		GLAND PHARMA LTD	<u>EQ 4MG PHOSPHATE/ML</u>	<u>A215654 001</u> Aug 04, 2021
<u>AP</u>			<u>EQ 10MG PHOSPHATE/ML</u>	<u>A215654 002</u> Sep 25, 2023
<u>AP</u>		HIKMA	<u>EQ 4MG PHOSPHATE/ML</u>	<u>A084282 001</u>
<u>AP</u>	!		<u>EQ 10MG PHOSPHATE/ML</u>	<u>A087702 001</u> Sep 07, 1982

PRESCRIPTION DRUG PRODUCT LIST

DEXAMETHASONE SODIUM PHOSPHATE

INJECTABLE; INJECTION

DEXAMETHASONE SODIUM PHOSPHATE

<u>AP</u>	MICRO LABS	<u>EQ 10MG PHOSPHATE/ML</u>	<u>A217797 001</u>	Mar 27, 2024
<u>AP</u>	MYLAN LABS LTD	<u>EQ 4MG PHOSPHATE/ML</u>	<u>A040803 001</u>	Aug 29, 2008
<u>AP</u>		<u>EQ 10MG PHOSPHATE/ML</u>	<u>A040802 001</u>	Aug 29, 2008
<u>AP</u>	SOMERSET	<u>EQ 4MG PHOSPHATE/ML</u>	<u>A207521 001</u>	Jun 08, 2018
<u>AP</u>	SOMERSET THERAPS LLC	<u>EQ 10MG PHOSPHATE/ML</u>	<u>A211036 001</u>	May 10, 2019

DEXAMETHASONE SODIUM PHOSPHATE PRESERVATIVE FREE

<u>AP</u>	AMNEAL	<u>EQ 10MG PHOSPHATE/ML</u>	<u>A208690 001</u>	Aug 22, 2018
<u>AP</u>	! FRESENIUS KABI USA	<u>EQ 10MG PHOSPHATE/ML</u>	<u>A040491 001</u>	Apr 11, 2003
<u>AP</u>	SOMERSET THERAPS LLC	<u>EQ 10MG PHOSPHATE/ML</u>	<u>A207442 001</u>	Apr 19, 2018

SOLUTION/DROPS; OPHTHALMIC, OTIC

DEXAMETHASONE SODIUM PHOSPHATE

!	BAUSCH AND LOMB	EQ 0.1% PHOSPHATE	A040069 001	Jul 26, 1996
---	-----------------	-------------------	-------------	--------------

DEXAMETHASONE; NEOMYCIN SULFATE; POLYMYXIN B SULFATE

OINTMENT; OPHTHALMIC

MAXITROL

<u>AT</u>	+! SANDOZ	<u>0.1%;EQ 3.5MG BASE/GM;10,000 UNITS/GM</u>	<u>N050065 002</u>	
<u>NEOMYCIN AND POLYMYXIN B SULFATES AND DEXAMETHASONE</u>				
<u>AT</u>	BAUSCH AND LOMB	<u>0.1%;EQ 3.5MG BASE/GM;10,000 UNITS/GM</u>	<u>A064063 001</u>	Jul 25, 1994
<u>AT</u>	PADAGIS US	<u>0.1%;EQ 3.5MG BASE/GM;10,000 UNITS/GM</u>	<u>A062938 001</u>	Jul 31, 1989

SUSPENSION/DROPS; OPHTHALMIC

DEXASPORIN

<u>AT</u>	BAUSCH AND LOMB	<u>0.1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML</u>	<u>A064135 001</u>	Sep 13, 1995
-----------	-----------------	--	--------------------	--------------

MAXITROL

<u>AT</u>	+! HARROW EYE	<u>0.1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML</u>	<u>N050023 002</u>	
<u>AT</u>	SANDOZ	<u>0.1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML</u>	<u>A062341 001</u>	May 22, 1984

DEXAMETHASONE; TOBRAMYCIN

OINTMENT; OPHTHALMIC

TOBRADEX

+	NOVARTIS	0.1%;0.3%	N050616 001	Sep 28, 1988
---	----------	-----------	-------------	--------------

SUSPENSION/DROPS; OPHTHALMIC

TOBRADEX

<u>AB</u>	+! SANDOZ	<u>0.1%;0.3%</u>	<u>N050592 001</u>	Aug 18, 1988
-----------	-----------	------------------	--------------------	--------------

TOBRAMYCIN AND DEXAMETHASONE

<u>AB</u>	BAUSCH AND LOMB	<u>0.1%;0.3%</u>	<u>A064134 001</u>	Oct 27, 1999
<u>AB</u>	PADAGIS US	<u>0.1%;0.3%</u>	<u>A212715 001</u>	Feb 11, 2022
	TOBRADEX ST			
+	HARROW EYE	0.05%;0.3%	N050818 001	Feb 13, 2009

DEXCHLORPHENIRAMINE MALEATE

SYRUP; ORAL

POLMON

!	CAPELLON PHARMS LLC	2MG/5ML	A202520 001	Jul 16, 2018
---	---------------------	---------	-------------	--------------

DEXLANSOPRAZOLE

CAPSULE, DELAYED RELEASE; ORAL

DEXILANT

<u>AB</u>	+ TAKEDA PHARMS USA	<u>30MG</u>	<u>N022287 001</u>	Jan 30, 2009
<u>AB</u>	+!	<u>60MG</u>	<u>N022287 002</u>	Jan 30, 2009

DEXLANSOPRAZOLE

<u>AB</u>	ENDO OPERATIONS	<u>30MG</u>	<u>A202294 002</u>	Jun 16, 2022
<u>AB</u>		<u>60MG</u>	<u>A202294 001</u>	Apr 19, 2017
<u>AB</u>	MYLAN	<u>30MG</u>	<u>A205205 001</u>	Jan 19, 2024
<u>AB</u>		<u>60MG</u>	<u>A205205 002</u>	Jan 19, 2024
<u>AB</u>	TWI PHARMS	<u>30MG</u>	<u>A202666 001</u>	Sep 16, 2022
<u>AB</u>		<u>60MG</u>	<u>A202666 002</u>	Sep 16, 2022

DEXMEDETOMIDINE HYDROCHLORIDE

FILM; BUCCAL, SUBLINGUAL

IGALMI

+	BIOXCEL	EQ 0.12MG BASE	N215390 001	Apr 05, 2022
+		EQ 0.18MG BASE	N215390 002	Apr 05, 2022

INJECTABLE; INJECTION

DEXMEDETOMIDINE HYDROCHLORIDE

<u>AP</u>	ACCORD HLTHCARE	<u>EQ 200MCG BASE/2ML (EQ 100MCG BASE/ML)</u>	<u>A204023 001</u>	Feb 09, 2016
<u>AP</u>	ACTAVIS INC	<u>EQ 200MCG BASE/2ML (EQ 100MCG BASE/ML)</u>	<u>A204686 001</u>	Oct 17, 2016
<u>AP</u>	AMNEAL	<u>EQ 400MCG BASE/50ML (EQ 4MCG BASE/ML)</u>	<u>A216604 001</u>	May 15, 2023
<u>AP</u>		<u>EQ 200MCG BASE/100ML (EQ 4MCG BASE/ML)</u>	<u>A216604 002</u>	May 15, 2023
<u>AP</u>	AMNEAL PHARMS CO	<u>EQ 200MCG BASE/50ML (EQ 4MCG BASE/ML)</u>	<u>A207551 001</u>	May 20, 2020

PRESCRIPTION DRUG PRODUCT LIST

DEXMEDETOMIDINE HYDROCHLORIDE

INJECTABLE; INJECTION

DEXMEDETOMIDINE HYDROCHLORIDE

<u>AP</u>		<u>EQ 400MCG BASE/100ML (EQ 4MCG BASE/ML)</u>	<u>A207551 002</u>	May 20, 2020
<u>AP</u>	BAXTER HLTHCARE CORP	<u>EQ 200MCG BASE/50ML (EQ 4MCG BASE/ML)</u>	<u>A208532 001</u>	Aug 21, 2018
<u>AP</u>		<u>EQ 400MCG BASE/100ML (EQ 4MCG BASE/ML)</u>	<u>A208532 002</u>	Aug 21, 2018
<u>AP</u>	ENDO OPERATIONS	<u>EQ 200MCG BASE/2ML (EQ 100MCG BASE/ML)</u>	<u>A203972 001</u>	Aug 18, 2014
<u>AP</u>		<u>EQ 200MCG BASE/50ML (EQ 4MCG BASE/ML)</u>	<u>A208266 001</u>	Sep 15, 2020
<u>AP</u>		<u>EQ 400MCG BASE/100ML (EQ 4MCG BASE/ML)</u>	<u>A208266 002</u>	Sep 15, 2020
<u>AP</u>	EUGIA PHARMA	<u>EQ 200MCG BASE/2ML (EQ 100MCG BASE/ML)</u>	<u>A205867 001</u>	Mar 17, 2016
<u>AP</u>		<u>EQ 200MCG BASE/50ML (EQ 4MCG BASE/ML)</u>	<u>A210321 001</u>	Dec 07, 2020
<u>AP</u>		<u>EQ 400MCG BASE/100ML (EQ 4MCG BASE/ML)</u>	<u>A210321 002</u>	Dec 07, 2020
<u>AP</u>	FRESENIUS KABI USA	<u>EQ 80MCG BASE/20ML (EQ 4MCG BASE/ML)</u>	<u>A208129 001</u>	Nov 29, 2018
<u>AP</u>		<u>EQ 200MCG BASE/2ML (EQ 100MCG BASE/ML)</u>	<u>A201072 001</u>	Sep 18, 2015
<u>AP</u>		<u>EQ 200MCG BASE/50ML (EQ 4MCG BASE/ML)</u>	<u>A208129 002</u>	Nov 29, 2018
<u>AP</u>		<u>EQ 400MCG BASE/100ML (EQ 4MCG BASE/ML)</u>	<u>A208129 003</u>	Nov 29, 2018
<u>AP</u>	GLAND	<u>EQ 200MCG BASE/2ML (EQ 100MCG BASE/ML)</u>	<u>A202126 001</u>	Aug 20, 2015
<u>AP</u>	GLAND PHARMA LTD	<u>EQ 200MCG BASE/50ML (EQ 4MCG BASE/ML)</u>	<u>A209307 001</u>	Jun 03, 2020
<u>AP</u>		<u>EQ 400MCG BASE/100ML (EQ 4MCG BASE/ML)</u>	<u>A209307 002</u>	Jun 03, 2020
<u>AP</u>	HENGRUI PHARMA	<u>EQ 80MCG BASE/20ML (EQ 4MCG BASE/ML)</u>	<u>A209065 002</u>	Jun 12, 2020
<u>AP</u>		<u>EQ 200MCG BASE/2ML (EQ 100MCG BASE/ML)</u>	<u>A209065 001</u>	Sep 19, 2017
<u>AP</u>		<u>EQ 200MCG BASE/50ML (EQ 4MCG BASE/ML)</u>	<u>A209065 003</u>	Jun 12, 2020
<u>AP</u>		<u>EQ 400MCG BASE/100ML (EQ 4MCG BASE/ML)</u>	<u>A209065 004</u>	Jun 12, 2020
<u>AP</u>	HIKMA	<u>EQ 200MCG BASE/2ML (EQ 100MCG BASE/ML)</u>	<u>A205046 001</u>	Apr 26, 2017
<u>AP</u>		<u>EQ 200MCG BASE/50ML (EQ 4MCG BASE/ML)</u>	<u>A206407 001</u>	Jan 30, 2020
<u>AP</u>		<u>EQ 400MCG BASE/100ML (EQ 4MCG BASE/ML)</u>	<u>A206407 002</u>	Jan 30, 2020
<u>AP</u>	MEITHEAL	<u>EQ 200MCG BASE/2ML (EQ 100MCG BASE/ML)</u>	<u>A204843 001</u>	Jan 18, 2019
<u>AP</u>	MILLA PHARMS	<u>EQ 200MCG BASE/50ML (EQ 4MCG BASE/ML)</u>	<u>A217308 001</u>	Jun 07, 2023
<u>AP</u>		<u>EQ 400MCG BASE/100ML (EQ 4MCG BASE/ML)</u>	<u>A217308 002</u>	Jun 07, 2023
<u>AP</u>	MYLAN INSTITUTIONAL	<u>EQ 200MCG BASE/2ML (EQ 100MCG BASE/ML)</u>	<u>A202881 001</u>	Aug 18, 2014
<u>AP</u>	MYLAN LABS LTD	<u>EQ 200MCG BASE/50ML (EQ 4MCG BASE/ML)</u>	<u>A212571 001</u>	Aug 27, 2020
<u>AP</u>		<u>EQ 400MCG BASE/100ML (EQ 4MCG BASE/ML)</u>	<u>A212571 002</u>	Aug 27, 2020
<u>AP</u>	RISING	<u>EQ 200MCG BASE/2ML (EQ 100MCG BASE/ML)</u>	<u>A202585 001</u>	Nov 24, 2014
<u>AP</u>	SANDOZ	<u>EQ 200MCG BASE/2ML (EQ 100MCG BASE/ML)</u>	<u>A091465 001</u>	Jun 14, 2016
<u>AP</u>	TAGI	<u>EQ 80MCG BASE/20ML (EQ 4MCG BASE/ML)</u>	<u>A212857 001</u>	Nov 23, 2020
<u>AP</u>		<u>EQ 200MCG BASE/50ML (EQ 4MCG BASE/ML)</u>	<u>A212857 002</u>	Nov 23, 2020
<u>AP</u>		<u>EQ 400MCG BASE/100ML (EQ 4MCG BASE/ML)</u>	<u>A212857 003</u>	Nov 23, 2020
<u>AP</u>	TEVA PHARMS USA	<u>EQ 200MCG BASE/2ML (EQ 100MCG BASE/ML)</u>	<u>A205272 001</u>	Nov 28, 2017
<u>AP</u>	WILSHIRE PHARMS INC	<u>EQ 200MCG BASE/50ML (EQ 4MCG BASE/ML)</u>	<u>A212791 001</u>	Dec 04, 2019
<u>AP</u>		<u>EQ 400MCG BASE/100ML (EQ 4MCG BASE/ML)</u>	<u>A212791 002</u>	Dec 04, 2019
<u>AP</u>	ZYDUS PHARMS	<u>EQ 200MCG BASE/2ML (EQ 100MCG BASE/ML)</u>	<u>A206798 001</u>	Feb 27, 2018

PRECEDEX

<u>AP</u>	+!	HOSPIRA	<u>EQ 80MCG BASE/20ML (EQ 4MCG BASE/ML)</u>	<u>N021038 004</u>	Nov 14, 2014
<u>AP</u>	+!		<u>EQ 200MCG BASE/2ML (EQ 100MCG BASE/ML)</u>	<u>N021038 001</u>	Dec 17, 1999
<u>AP</u>	+!		<u>EQ 200MCG BASE/50ML (EQ 4MCG BASE/ML)</u>	<u>N021038 002</u>	Mar 13, 2013
<u>AP</u>	+!		<u>EQ 400MCG BASE/100ML (EQ 4MCG BASE/ML)</u>	<u>N021038 003</u>	Mar 13, 2013
			EQ 1MG BASE/250ML (EQ 4MCG BASE/ML)	N021038 005	Jan 31, 2020

SOLUTION; INTRAVENOUS

DEXMEDETOMIDINE HYDROCHLORIDE

<u>AP</u>	+!	HQ SPCLT PHARMA	<u>EQ 1MG BASE/10ML (EQ 100MCG BASE/ML)</u>	<u>N206628 002</u>	Oct 21, 2015
<u>AP</u>	+		<u>EQ 400MCG BASE/4ML (EQ 100MCG BASE/ML)</u>	<u>N206628 001</u>	Oct 21, 2015
<u>AP</u>		SOMERSET	<u>EQ 1MG BASE/10ML (EQ 100MCG BASE/ML)</u>	<u>A218112 002</u>	Sep 24, 2024
<u>AP</u>			<u>EQ 400MCG BASE/4ML (EQ 100MCG BASE/ML)</u>	<u>A218112 001</u>	Sep 24, 2024
	+	HQ SPCLT PHARMA	EQ 200MCG BASE/50ML (EQ 4MCG BASE/ML)	N206628 003	Jun 22, 2018
	+		EQ 400MCG BASE/100ML (EQ 4MCG BASE/ML)	N206628 004	Jun 22, 2018

DEXMETHYLPHENIDATE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

DEXMETHYLPHENIDATE HYDROCHLORIDE

<u>AB</u>		ADARE PHARMS INC	<u>5MG</u>	<u>A210279 001</u>	Oct 09, 2018
<u>AB</u>			<u>10MG</u>	<u>A210279 002</u>	Oct 09, 2018
<u>AB</u>			<u>15MG</u>	<u>A210279 003</u>	Oct 09, 2018
<u>AB</u>			<u>20MG</u>	<u>A210279 004</u>	Oct 09, 2018
<u>AB</u>			<u>25MG</u>	<u>A210279 005</u>	Oct 09, 2018
<u>AB</u>			<u>30MG</u>	<u>A210279 006</u>	Oct 09, 2018
<u>AB</u>			<u>35MG</u>	<u>A210279 007</u>	Oct 09, 2018
<u>AB</u>			<u>40MG</u>	<u>A210279 008</u>	Oct 09, 2018
<u>AB</u>		ASCENT PHARMS INC	<u>5MG</u>	<u>A215523 001</u>	Dec 08, 2021
<u>AB</u>			<u>10MG</u>	<u>A215523 002</u>	Dec 08, 2021
<u>AB</u>			<u>15MG</u>	<u>A215523 003</u>	Dec 08, 2021
<u>AB</u>			<u>20MG</u>	<u>A215523 004</u>	Dec 08, 2021
<u>AB</u>			<u>25MG</u>	<u>A215523 005</u>	Dec 08, 2021

PRESCRIPTION DRUG PRODUCT LIST

DEXMETHYLPHENIDATE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE;ORAL

DEXMETHYLPHENIDATE HYDROCHLORIDE

<u>AB</u>		<u>30MG</u>	<u>A215523 006</u>	Dec 08, 2021
<u>AB</u>		<u>35MG</u>	<u>A215523 007</u>	Dec 08, 2021
<u>AB</u>		<u>40MG</u>	<u>A215523 008</u>	Dec 08, 2021
<u>AB</u>	ENDO OPERATIONS	<u>5MG</u>	<u>A202842 001</u>	Nov 30, 2016
<u>AB</u>		<u>10MG</u>	<u>A202842 002</u>	Nov 30, 2016
<u>AB</u>		<u>15MG</u>	<u>A202842 003</u>	Nov 30, 2016
<u>AB</u>		<u>20MG</u>	<u>A202842 004</u>	Nov 30, 2016
<u>AB</u>		<u>25MG</u>	<u>A202842 005</u>	Nov 30, 2016
<u>AB</u>		<u>30MG</u>	<u>A202842 006</u>	Nov 30, 2016
<u>AB</u>		<u>35MG</u>	<u>A202842 007</u>	Nov 30, 2016
<u>AB</u>		<u>40MG</u>	<u>A202842 008</u>	Nov 30, 2016
<u>AB</u>	GRANULES	<u>5MG</u>	<u>A213813 001</u>	Sep 09, 2020
<u>AB</u>		<u>10MG</u>	<u>A213813 002</u>	Sep 09, 2020
<u>AB</u>		<u>15MG</u>	<u>A213813 003</u>	Sep 09, 2020
<u>AB</u>		<u>20MG</u>	<u>A213813 004</u>	Sep 09, 2020
<u>AB</u>		<u>25MG</u>	<u>A213813 005</u>	Sep 09, 2020
<u>AB</u>		<u>30MG</u>	<u>A213813 006</u>	Sep 09, 2020
<u>AB</u>		<u>35MG</u>	<u>A213813 007</u>	Sep 09, 2020
<u>AB</u>		<u>40MG</u>	<u>A213813 008</u>	Sep 09, 2020
<u>AB</u>	IMPAX LABS INC	<u>5MG</u>	<u>A079108 001</u>	Aug 05, 2015
<u>AB</u>		<u>10MG</u>	<u>A079108 002</u>	Aug 05, 2015
<u>AB</u>		<u>15MG</u>	<u>A079108 003</u>	May 19, 2014
<u>AB</u>		<u>20MG</u>	<u>A079108 004</u>	Dec 21, 2015
<u>AB</u>		<u>25MG</u>	<u>A203614 001</u>	Jul 05, 2017
<u>AB</u>		<u>30MG</u>	<u>A079108 005</u>	Nov 21, 2013
<u>AB</u>		<u>35MG</u>	<u>A203614 002</u>	Jul 05, 2017
<u>AB</u>	INTELLIPHARMACEUTIC S	<u>5MG</u>	<u>A078992 001</u>	Nov 23, 2021
<u>AB</u>		<u>10MG</u>	<u>A078992 002</u>	Nov 23, 2021
<u>AB</u>		<u>15MG</u>	<u>A078992 003</u>	Nov 18, 2013
<u>AB</u>		<u>30MG</u>	<u>A078992 004</u>	Nov 18, 2013
<u>AB</u>		<u>20MG</u>	<u>A078992 005</u>	Nov 23, 2021
<u>AB</u>		<u>40MG</u>	<u>A078992 006</u>	Nov 23, 2021
<u>AB</u>	SUN PHARM INDS INC	<u>5MG</u>	<u>A206734 001</u>	Nov 05, 2021
<u>AB</u>		<u>10MG</u>	<u>A206734 002</u>	Nov 05, 2021
<u>AB</u>		<u>15MG</u>	<u>A206734 003</u>	Nov 05, 2021
<u>AB</u>		<u>20MG</u>	<u>A206734 004</u>	Nov 05, 2021
<u>AB</u>		<u>25MG</u>	<u>A206734 005</u>	Nov 05, 2021
<u>AB</u>		<u>30MG</u>	<u>A206734 006</u>	Nov 05, 2021
<u>AB</u>		<u>35MG</u>	<u>A206734 007</u>	Nov 05, 2021
<u>AB</u>		<u>40MG</u>	<u>A206734 008</u>	Nov 05, 2021
<u>AB</u>	TEVA PHARMS USA	<u>5MG</u>	<u>A078908 001</u>	Nov 19, 2013
<u>AB</u>		<u>10MG</u>	<u>A078908 002</u>	Nov 19, 2013
<u>AB</u>		<u>15MG</u>	<u>A078908 004</u>	May 19, 2014
<u>AB</u>		<u>20MG</u>	<u>A078908 003</u>	Nov 19, 2013
<u>AB</u>		<u>25MG</u>	<u>A202731 001</u>	Jul 05, 2017
<u>AB</u>		<u>30MG</u>	<u>A202731 003</u>	May 19, 2014
<u>AB</u>		<u>35MG</u>	<u>A202731 004</u>	Jul 05, 2017
<u>AB</u>		<u>40MG</u>	<u>A202731 002</u>	Nov 19, 2013

FOCALIN XR

<u>AB</u>	+	SANDOZ	<u>5MG</u>	<u>N021802 001</u>	May 26, 2005
<u>AB</u>	+		<u>10MG</u>	<u>N021802 002</u>	May 26, 2005
<u>AB</u>	+		<u>15MG</u>	<u>N021802 004</u>	Aug 01, 2006
<u>AB</u>	+		<u>20MG</u>	<u>N021802 003</u>	May 26, 2005
<u>AB</u>	+		<u>25MG</u>	<u>N021802 008</u>	Apr 21, 2011
<u>AB</u>	+		<u>30MG</u>	<u>N021802 005</u>	Oct 23, 2009
<u>AB</u>	+		<u>35MG</u>	<u>N021802 007</u>	Apr 21, 2011
<u>AB</u>	+		<u>40MG</u>	<u>N021802 006</u>	Aug 11, 2010

TABLET;ORAL

DEXMETHYLPHENIDATE HYDROCHLORIDE

<u>AB</u>	ABHAI INC	<u>2.5MG</u>	<u>A206931 001</u>	Dec 04, 2015
<u>AB</u>		<u>5MG</u>	<u>A206931 002</u>	Dec 04, 2015
<u>AB</u>		<u>10MG</u>	<u>A206931 003</u>	Dec 04, 2015
<u>AB</u>	ALKEM LABS LTD	<u>2.5MG</u>	<u>A212631 001</u>	Jul 19, 2019
<u>AB</u>		<u>5MG</u>	<u>A212631 002</u>	Jul 19, 2019
<u>AB</u>		<u>10MG</u>	<u>A212631 003</u>	Jul 19, 2019
<u>AB</u>	CEDIPROF INC	<u>5MG</u>	<u>A209211 001</u>	Sep 19, 2018
<u>AB</u>		<u>10MG</u>	<u>A209211 002</u>	Sep 19, 2018
<u>AB</u>	NOVEL LABS INC	<u>2.5MG</u>	<u>A204534 001</u>	Dec 04, 2015
<u>AB</u>		<u>5MG</u>	<u>A204534 002</u>	Dec 04, 2015

PRESCRIPTION DRUG PRODUCT LIST

DEXMETHYLPHENIDATE HYDROCHLORIDE

TABLET; ORAL

DEXMETHYLPHENIDATE HYDROCHLORIDE

<u>AB</u>		<u>10MG</u>	<u>A204534</u>	<u>003</u>	Dec 04, 2015
<u>AB</u>	RHODES PHARMS	<u>2.5MG</u>	<u>A208756</u>	<u>001</u>	Nov 20, 2017
<u>AB</u>		<u>5MG</u>	<u>A208756</u>	<u>002</u>	Nov 20, 2017
<u>AB</u>		<u>10MG</u>	<u>A208756</u>	<u>003</u>	Nov 20, 2017
<u>AB</u>	SUN PHARM INDUSTRIES	<u>2.5MG</u>	<u>A201231</u>	<u>001</u>	Sep 24, 2015
<u>AB</u>		<u>5MG</u>	<u>A201231</u>	<u>002</u>	Sep 24, 2015
<u>AB</u>		<u>10MG</u>	<u>A201231</u>	<u>003</u>	Sep 24, 2015
<u>AB</u>	TRIS PHARMA INC	<u>2.5MG</u>	<u>A207901</u>	<u>001</u>	Aug 26, 2016
<u>AB</u>		<u>5MG</u>	<u>A207901</u>	<u>002</u>	Aug 26, 2016
<u>AB</u>		<u>10MG</u>	<u>A207901</u>	<u>003</u>	Aug 26, 2016
<u>FOCALIN</u>					
<u>AB</u>	+ SANDOZ	<u>2.5MG</u>	<u>N021278</u>	<u>001</u>	Nov 13, 2001
<u>AB</u>	+	<u>5MG</u>	<u>N021278</u>	<u>002</u>	Nov 13, 2001
<u>AB</u>	+!	<u>10MG</u>	<u>N021278</u>	<u>003</u>	Nov 13, 2001

DEXMETHYLPHENIDATE HYDROCHLORIDE; SERDEXMETHYLPHENIDATE CHLORIDE

CAPSULE; ORAL

AZSTARYS

+	COMMAVE THERAP	EQ 5.2MG BASE;EQ 26.1MG BASE	N212994	001	May 07, 2021
+		EQ 7.8MG BASE;EQ 39.2MG BASE	N212994	002	May 07, 2021
+!		EQ 10.4MG BASE;EQ 52.3MG BASE	N212994	003	May 07, 2021

DEXRAZOXANE HYDROCHLORIDE

INJECTABLE; INJECTION

DEXRAZOXANE HYDROCHLORIDE

<u>AP</u>	!	EUGIA PHARMA	<u>EQ 250MG BASE/VIAL</u>	<u>A200752</u>	<u>001</u>	Oct 19, 2011
<u>AP</u>	!		<u>EQ 500MG BASE/VIAL</u>	<u>A200752</u>	<u>002</u>	Oct 19, 2011
<u>AP</u>		GLAND	<u>EQ 250MG BASE/VIAL</u>	<u>A207321</u>	<u>002</u>	Dec 16, 2019
<u>AP</u>			<u>EQ 500MG BASE/VIAL</u>	<u>A207321</u>	<u>001</u>	Nov 28, 2016
<u>AP</u>		HIKMA	<u>EQ 250MG BASE/VIAL</u>	<u>A076068</u>	<u>001</u>	Sep 28, 2004
<u>AP</u>			<u>EQ 500MG BASE/VIAL</u>	<u>A076068</u>	<u>002</u>	Sep 28, 2004

DEXTROAMPHETAMINE

SYSTEM; TRANSDERMAL

KELSTRYM

+	NOVEN PHARMS INC	4.5MG/9HR	N215401	001	Mar 22, 2022
+		9MG/9HR	N215401	002	Mar 22, 2022
+		13.5MG/9HR	N215401	003	Mar 22, 2022
+!		18MG/9HR	N215401	004	Mar 22, 2022

DEXTROAMPHETAMINE SULFATE

CAPSULE, EXTENDED RELEASE; ORAL

DEXEDRINE SPANSULE

<u>AB</u>	+	IMPAX LABS INC	<u>5MG</u>	<u>N017078</u>	<u>001</u>
<u>AB</u>	+		<u>10MG</u>	<u>N017078</u>	<u>002</u>
<u>AB</u>	+!		<u>15MG</u>	<u>N017078</u>	<u>003</u>

DEXTROAMPHETAMINE SULFATE

<u>AB</u>		ACTAVIS ELIZABETH	<u>5MG</u>	<u>A203901</u>	<u>001</u>	Nov 30, 2012
<u>AB</u>			<u>10MG</u>	<u>A203901</u>	<u>002</u>	Nov 30, 2012
<u>AB</u>			<u>15MG</u>	<u>A203901</u>	<u>003</u>	Nov 30, 2012
<u>AB</u>		SPECGX LLC	<u>5MG</u>	<u>A076353</u>	<u>001</u>	May 06, 2003
<u>AB</u>			<u>10MG</u>	<u>A076353</u>	<u>002</u>	May 06, 2003
<u>AB</u>			<u>15MG</u>	<u>A076353</u>	<u>003</u>	May 06, 2003
<u>AB</u>		STRIDES PHARMA	<u>5MG</u>	<u>A205673</u>	<u>001</u>	Oct 31, 2017
<u>AB</u>			<u>10MG</u>	<u>A205673</u>	<u>002</u>	Oct 31, 2017
<u>AB</u>			<u>15MG</u>	<u>A205673</u>	<u>003</u>	Oct 31, 2017

SOLUTION; ORAL

DEXTROAMPHETAMINE SULFATE

<u>AA</u>	!	PRASCO	<u>5MG/5ML</u>	<u>A040776</u>	<u>001</u>	Jan 29, 2008
<u>AA</u>		TRIS PHARMA INC	<u>5MG/5ML</u>	<u>A203644</u>	<u>001</u>	May 29, 2013

TABLET; ORAL

DEXTROAMPHETAMINE SULFATE

<u>AA</u>		AUROLIFE PHARMA LLC	<u>5MG</u>	<u>A202893</u>	<u>001</u>	Jul 31, 2013
<u>AA</u>			<u>10MG</u>	<u>A202893</u>	<u>002</u>	Jul 31, 2013
<u>AA</u>		AVANTHI INC	<u>5MG</u>	<u>A203548</u>	<u>001</u>	Nov 23, 2015
<u>AA</u>			<u>10MG</u>	<u>A203548</u>	<u>002</u>	Nov 23, 2015
<u>AA</u>		AZURITY	<u>2.5MG</u>	<u>A090533</u>	<u>001</u>	Oct 25, 2011
<u>AA</u>			<u>5MG</u>	<u>A090533</u>	<u>002</u>	Oct 25, 2011
<u>AA</u>			<u>7.5MG</u>	<u>A090533</u>	<u>003</u>	Oct 25, 2011
<u>AA</u>			<u>10MG</u>	<u>A090533</u>	<u>004</u>	Oct 25, 2011
<u>AA</u>			<u>15MG</u>	<u>A090533</u>	<u>005</u>	Oct 25, 2011

PRESCRIPTION DRUG PRODUCT LIST

DEXTROAMPHETAMINE SULFATE

TABLET; ORAL

DEXTROAMPHETAMINE SULFATE

<u>AA</u>		<u>20MG</u>	<u>A090533 006</u>	Oct 25, 2011
<u>AA</u>		<u>30MG</u>	<u>A090533 007</u>	Oct 25, 2011
<u>AA</u>	BARR	<u>5MG</u>	<u>A040361 001</u>	Jan 31, 2001
<u>AA</u>	!	<u>10MG</u>	<u>A040361 002</u>	Jan 31, 2001
<u>AA</u>	NOVEL LABS INC	<u>5MG</u>	<u>A204330 001</u>	Mar 16, 2016
<u>AA</u>		<u>10MG</u>	<u>A204330 002</u>	Mar 16, 2016
<u>AA</u>	NUVO PHARM	<u>5MG</u>	<u>A210059 001</u>	Oct 18, 2017
<u>AA</u>		<u>10MG</u>	<u>A210059 002</u>	Oct 18, 2017
<u>AA</u>	SPECGX LLC	<u>5MG</u>	<u>A040436 001</u>	Jan 29, 2002
<u>AA</u>		<u>10MG</u>	<u>A040436 002</u>	Jan 29, 2002
<u>AA</u>	WINDER LABS LLC	<u>2.5MG</u>	<u>A212160 001</u>	Jun 07, 2021
<u>AA</u>		<u>5MG</u>	<u>A212160 002</u>	Jun 07, 2021
<u>AA</u>		<u>7.5MG</u>	<u>A212160 003</u>	Jun 07, 2021
<u>AA</u>		<u>10MG</u>	<u>A212160 004</u>	Jun 07, 2021
<u>AA</u>		<u>15MG</u>	<u>A212160 005</u>	Jun 07, 2021
<u>AA</u>		<u>20MG</u>	<u>A212160 006</u>	Jun 07, 2021
<u>AA</u>		<u>30MG</u>	<u>A212160 007</u>	Jun 07, 2021

DEXTROMETHORPHAN HYDROBROMIDE; PROMETHAZINE HYDROCHLORIDE

SYRUP; ORAL

PROMETHAZINE DM

<u>AA</u>	!	SLATE RUN PHARMA	<u>15MG/5ML; 6.25MG/5ML</u>	<u>A040649 001</u>	Feb 14, 2006
<u>PROMETHAZINE HYDROCHLORIDE AND DEXTROMETHORPHAN HYDROBROMIDE</u>					
<u>AA</u>		AMNEAL PHARMS	<u>15MG/5ML; 6.25MG/5ML</u>	<u>A090575 001</u>	Feb 08, 2011
<u>AA</u>	+	ANI PHARMS	<u>15MG/5ML; 6.25MG/5ML</u>	<u>N011265 002</u>	Apr 02, 1984
<u>AA</u>		TRIS PHARMA INC	<u>15MG/5ML; 6.25MG/5ML</u>	<u>A091687 001</u>	Jun 28, 2012
<u>PROMETHAZINE W/ DEXTROMETHORPHAN</u>					
<u>AA</u>		WOCKHARDT BIO AG	<u>15MG/5ML; 6.25MG/5ML</u>	<u>A088864 001</u>	Jan 04, 1985

DEXTROMETHORPHAN HYDROBROMIDE; QUINIDINE SULFATE

CAPSULE; ORAL

DEXTROMETHORPHAN HYDROBROMIDE AND QUINIDINE SULFATE

<u>AB</u>		ACTAVIS ELIZABETH	<u>20MG; 10MG</u>	<u>A202934 001</u>	Oct 10, 2017
<u>AB</u>		HETERO LABS LTD III	<u>20MG; 10MG</u>	<u>A218426 001</u>	Aug 28, 2024
<u>NUDEXTA</u>					
<u>AB</u>	+	AVANIR PHARMS	<u>20MG; 10MG</u>	<u>N021879 001</u>	Oct 29, 2010

DEXTROSE

INJECTABLE; INJECTION

DEXTROSE 10% IN PLASTIC CONTAINER

<u>AP</u>	+	B BRAUN	<u>10GM/100ML</u>	<u>N019626 004</u>	Feb 02, 1988
<u>AP</u>	+	BAXTER HLTHCARE	<u>10GM/100ML</u>	<u>N016694 001</u>	
<u>AP</u>		FRESENIUS KABI USA	<u>10GM/100ML</u>	<u>A209448 001</u>	Jul 16, 2018
<u>AP</u>	+	ICU MEDICAL INC	<u>10GM/100ML</u>	<u>N018080 001</u>	

DEXTROSE 5% IN PLASTIC CONTAINER

<u>AP</u>	+	B BRAUN	<u>50MG/ML</u>	<u>N016730 002</u>	
<u>AP</u>	+		<u>5GM/100ML</u>	<u>N016730 001</u>	
<u>AP</u>	+		<u>5GM/100ML</u>	<u>N019626 002</u>	Feb 02, 1988
<u>AP</u>	+	BAXTER HLTHCARE	<u>50MG/ML</u>	<u>N016673 003</u>	Oct 30, 1985
<u>AP</u>	+		<u>50MG/ML</u>	<u>N020179 002</u>	Dec 07, 1992
<u>AP</u>	+		<u>5GM/100ML</u>	<u>N016673 001</u>	
<u>AP</u>	+		<u>5GM/100ML</u>	<u>N020179 001</u>	Dec 07, 1992
<u>AP</u>		FRESENIUS KABI USA	<u>50MG/ML</u>	<u>A207449 001</u>	Oct 21, 2016
<u>AP</u>	+	HOSPIRA	<u>5GM/100ML</u>	<u>N019466 001</u>	Jul 15, 1985
<u>AP</u>	+		<u>5GM/100ML</u>	<u>N019479 001</u>	Sep 17, 1985
<u>AP</u>	+	ICU MEDICAL INC	<u>50MG/ML</u>	<u>N016367 002</u>	

DEXTROSE 50%

<u>AP</u>	+	HOSPIRA	<u>500MG/ML</u>	<u>N019445 001</u>	Jun 03, 1986
<u>AP</u>		INTL MEDICATION SYS	<u>500MG/ML</u>	<u>A203451 001</u>	Mar 26, 2021

DEXTROSE 50% IN PLASTIC CONTAINER

<u>AP</u>	+	BAXTER HLTHCARE	<u>50GM/100ML</u>	<u>N020047 001</u>	Jul 02, 1991
<u>AP</u>	+	ICU MEDICAL INC	<u>50GM/100ML</u>	<u>N018563 001</u>	Mar 23, 1982

DEXTROSE 70% IN PLASTIC CONTAINER

<u>AP</u>	+	B BRAUN	<u>70GM/100ML</u>	<u>N019626 005</u>	Feb 18, 2015
<u>AP</u>	+	BAXTER HLTHCARE	<u>70GM/100ML</u>	<u>N020047 003</u>	Jul 02, 1991
<u>AP</u>	+	ICU MEDICAL INC	<u>70GM/100ML</u>	<u>N018561 001</u>	Mar 23, 1982
<u>AP</u>	+		<u>70GM/100ML</u>	<u>N019893 001</u>	Dec 26, 1989

DEXTROSE 20% IN PLASTIC CONTAINER

	+	ICU MEDICAL INC	<u>20GM/100ML</u>	<u>N018564 001</u>	Mar 23, 1982
--	---	-----------------	-------------------	--------------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

DEXTROSE

INJECTABLE; INJECTION

DEXTROSE 25%

+! HOSPIRA 250MG/ML N019445 002 Nov 23, 1998

DEXTROSE 30% IN PLASTIC CONTAINER

+! ICU MEDICAL INC 30GM/100ML N019345 001 Jan 26, 1985

DEXTROSE 40% IN PLASTIC CONTAINER

+! ICU MEDICAL INC 40GM/100ML N018562 001 Mar 23, 1982

DEXTROSE; MAGNESIUM ACETATE; POTASSIUM ACETATE; SODIUM CHLORIDE

INJECTABLE; INJECTION

NORMOSOL-M AND DEXTROSE 5% IN PLASTIC CONTAINER

ICU MEDICAL INC 5GM/100ML; 21MG/100ML; 128MG/100ML; 234MG/100ML N017610 001

DEXTROSE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; POTASSIUM PHOSPHATE, DIBASIC; SODIUM ACETATE

INJECTABLE; INJECTION

ISOLYTE P IN DEXTROSE 5% IN PLASTIC CONTAINER

B BRAUN 5GM/100ML; 31MG/100ML; 130MG/100ML; 26MG/100ML; 320MG/100ML N019873 001 Jun 10, 1993

DEXTROSE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; POTASSIUM PHOSPHATE, MONOBASIC; SODIUM CHLORIDE; SODIUM LACTATE

INJECTABLE; INJECTION

DEXTROSE 5% AND ELECTROLYTE NO. 48 IN PLASTIC CONTAINER

BAXTER HLTHCARE 5GM/100ML; 31MG/100ML; 141MG/100ML; 20MG/100ML; 12MG/100ML; 260MG/100ML N017484 001

DEXTROSE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; POTASSIUM PHOSPHATE, MONOBASIC; SODIUM LACTATE; SODIUM PHOSPHATE, MONOBASIC ANHYDROUS

INJECTABLE; INJECTION

IONOSOL MB AND DEXTROSE 5% IN PLASTIC CONTAINER

ICU MEDICAL INC 5GM/100ML; 30MG/100ML; 141MG/100ML; 15MG/100ML; 260MG/100ML; 25MG/100ML N019513 001 May 08, 1986

DEXTROSE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE; SODIUM GLUCONATE

INJECTABLE; INJECTION

NORMOSOL-R AND DEXTROSE 5% IN PLASTIC CONTAINER

ICU MEDICAL INC 5GM/100ML; 30MG/100ML; 37MG/100ML; 222MG/100ML; 526MG/100ML; 502MG/100ML N017609 001

DEXTROSE; POTASSIUM CHLORIDE

INJECTABLE; INJECTION

DEXTROSE 5% AND POTASSIUM CHLORIDE 0.075%AP FRESENIUS KABI USA 5GM/100ML; 75MG/100ML A212346 001 Sep 10, 2020DEXTROSE 5% AND POTASSIUM CHLORIDE 0.075% IN PLASTIC CONTAINERAP + BAXTER HLTHCARE 5GM/100ML; 75MG/100ML N017634 004DEXTROSE 5% AND POTASSIUM CHLORIDE 0.15%AP FRESENIUS KABI USA 5GM/100ML; 150MG/100ML A212346 002 Sep 10, 2020DEXTROSE 5% AND POTASSIUM CHLORIDE 0.15% IN PLASTIC CONTAINERAP + BAXTER HLTHCARE 5GM/100ML; 150MG/100ML N017634 001DEXTROSE 5% AND POTASSIUM CHLORIDE 0.3% IN PLASTIC CONTAINERAP + BAXTER HLTHCARE 5GM/100ML; 300MG/100ML N017634 002POTASSIUM CHLORIDE 0.15% IN DEXTROSE 5% IN PLASTIC CONTAINERAP B BRAUN 5GM/100ML; 150MG/100ML N019699 004 Sep 29, 1989POTASSIUM CHLORIDE 0.3% IN DEXTROSE 5% IN PLASTIC CONTAINERAP B BRAUN 5GM/100ML; 300MG/100ML N019699 006 Sep 29, 1989

DEXTROSE 5% AND POTASSIUM CHLORIDE 0.224% IN PLASTIC CONTAINER

+ BAXTER HLTHCARE 5GM/100ML; 224MG/100ML N017634 003

POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% IN PLASTIC CONTAINER

ICU MEDICAL INC 5GM/100ML; 149MG/100ML N018371 001

DEXTROSE; POTASSIUM CHLORIDE; SODIUM CHLORIDE

INJECTABLE; INJECTION

DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 10MEQAP BAXTER HLTHCARE 5GM/100ML; 75MG/100ML; 200MG/100ML N018037 006 Apr 13, 1982AP 5GM/100ML; 150MG/100ML; 200MG/100ML N018037 007 Apr 13, 1982DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 15MEQ (K)AP BAXTER HLTHCARE 5GM/100ML; 224MG/100ML; 200MG/100ML N018037 004DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 20MEQAP BAXTER HLTHCARE 5GM/100ML; 150MG/100ML; 200MG/100ML N018037 008 Apr 13, 1982DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 20MEQ (K)AP BAXTER HLTHCARE 5GM/100ML; 300MG/100ML; 200MG/100ML N018037 001DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 30MEQAP BAXTER HLTHCARE 5GM/100ML; 224MG/100ML; 200MG/100ML N018037 005 Apr 13, 1982

PRESCRIPTION DRUG PRODUCT LIST

DEXTROSE; POTASSIUM CHLORIDE; SODIUM CHLORIDEINJECTABLE; INJECTION

		<u>DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 40MEQ</u>		
AP	BAXTER HLTHCARE	5GM/100ML;300MG/100ML;200MG/100ML	N018037 009	Apr 13, 1982
		<u>DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 5MEQ</u>		
AP	BAXTER HLTHCARE	5GM/100ML;75MG/100ML;200MG/100ML	N018037 002	
		<u>DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 5MEQ (K)</u>		
AP	BAXTER HLTHCARE	5GM/100ML;150MG/100ML;200MG/100ML	N018037 003	
		<u>DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 10MEQ IN PLASTIC CONTAINER</u>		
AP	BAXTER HLTHCARE	5GM/100ML;75MG/100ML;330MG/100ML	N018629 005	Mar 23, 1982
AP		5GM/100ML;150MG/100ML;330MG/100ML	N018629 002	Mar 23, 1982
		<u>DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 15MEQ IN PLASTIC CONTAINER</u>		
AP	BAXTER HLTHCARE	5GM/100ML;224MG/100ML;330MG/100ML	N018629 003	Mar 23, 1982
		<u>DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 20MEQ IN PLASTIC CONTAINER</u>		
AP	BAXTER HLTHCARE	5GM/100ML;150MG/100ML;330MG/100ML	N018629 004	Mar 23, 1982
AP		5GM/100ML;300MG/100ML;330MG/100ML	N018629 006	Mar 23, 1982
		<u>DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 30MEQ IN PLASTIC CONTAINER</u>		
AP	BAXTER HLTHCARE	5GM/100ML;224MG/100ML;330MG/100ML	N018629 007	Mar 23, 1982
		<u>DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 40MEQ IN PLASTIC CONTAINER</u>		
AP	BAXTER HLTHCARE	5GM/100ML;300MG/100ML;330MG/100ML	N018629 008	Mar 23, 1982
		<u>DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 5MEQ IN PLASTIC CONTAINER</u>		
AP	BAXTER HLTHCARE	5GM/100ML;75MG/100ML;330MG/100ML	N018629 001	Mar 23, 1982
		<u>DEXTROSE 5%, SODIUM CHLORIDE 0.45% AND POTASSIUM CHLORIDE 20MEQ (K) IN PLASTIC CONTAINER</u>		
AP	BAXTER HLTHCARE	5GM/100ML;300MG/100ML;450MG/100ML	N018008 010	
		<u>POTASSIUM CHLORIDE 0.075% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER</u>		
AP	B BRAUN	5GM/100ML;75MG/100ML;200MG/100ML	N019630 008	Feb 17, 1988
		<u>POTASSIUM CHLORIDE 0.075% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER</u>		
AP	B BRAUN	5GM/100ML;75MG/100ML;330MG/100ML	N019630 014	Feb 17, 1988
		<u>POTASSIUM CHLORIDE 0.075% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER</u>		
AP	B BRAUN	5GM/100ML;75MG/100ML;450MG/100ML	N019630 020	Feb 17, 1988
		<u>POTASSIUM CHLORIDE 0.075% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER</u>		
AP	B BRAUN	5GM/100ML;75MG/100ML;900MG/100ML	N019630 026	Feb 17, 1988
		<u>POTASSIUM CHLORIDE 0.15% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER</u>		
AP	B BRAUN	5GM/100ML;150MG/100ML;200MG/100ML	N019630 010	Feb 17, 1988
		<u>POTASSIUM CHLORIDE 0.15% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER</u>		
AP	B BRAUN	5GM/100ML;150MG/100ML;330MG/100ML	N019630 016	Feb 17, 1988
		<u>POTASSIUM CHLORIDE 0.15% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER</u>		
AP	B BRAUN	5GM/100ML;150MG/100ML;450MG/100ML	N019630 022	Feb 17, 1988
		<u>POTASSIUM CHLORIDE 0.15% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER</u>		
AP	B BRAUN	5GM/100ML;150MG/100ML;900MG/100ML	N019630 028	Feb 17, 1988
		<u>POTASSIUM CHLORIDE 0.3% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER</u>		
AP	B BRAUN	5GM/100ML;300MG/100ML;200MG/100ML	N019630 012	Feb 17, 1988
		<u>POTASSIUM CHLORIDE 0.3% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER</u>		
AP	B BRAUN	5GM/100ML;300MG/100ML;330MG/100ML	N019630 018	Feb 17, 1988
		<u>POTASSIUM CHLORIDE 0.3% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER</u>		
AP	B BRAUN	5GM/100ML;300MG/100ML;450MG/100ML	N019630 024	Feb 17, 1988
		<u>POTASSIUM CHLORIDE 0.3% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER</u>		
AP	B BRAUN	5GM/100ML;300MG/100ML;900MG/100ML	N019630 030	Feb 17, 1988
		<u>POTASSIUM CHLORIDE 10MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.225% IN PLASTIC CONTAINER</u>		
AP	FRESENIUS KABI USA	5GM/100ML;149MG/100ML;225MG/100ML	A212348 001	Jul 30, 2021
		<u>POTASSIUM CHLORIDE 10MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45%</u>		
AP	FRESENIUS KABI USA	5GM/100ML;74.5MG/100ML;450MG/100ML	A213523 001	Mar 09, 2021
AP		5GM/100ML;149MG/100ML;450MG/100ML	A213523 005	Oct 11, 2022
		<u>POTASSIUM CHLORIDE 10MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER</u>		
AP	BAXTER HLTHCARE	5GM/100ML;75MG/100ML;450MG/100ML	N018008 005	Apr 28, 1982
AP		5GM/100ML;150MG/100ML;450MG/100ML	N018008 006	Apr 28, 1982
AP	+! ICU MEDICAL INC	5GM/100ML;74.5MG/100ML;450MG/100ML	N018362 009	Jul 05, 1983
AP	+!	5GM/100ML;149MG/100ML;450MG/100ML	N018362 005	Mar 28, 1988
		<u>POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.225% IN PLASTIC CONTAINER</u>		
AP	FRESENIUS KABI USA	5GM/100ML;149MG/100ML;225MG/100ML	A212348 002	Jul 30, 2021
AP	+! ICU MEDICAL INC	5GM/100ML;149MG/100ML;225MG/100ML	N018365 001	
		<u>POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45%</u>		
AP	FRESENIUS KABI USA	5GM/100ML;149MG/100ML;450MG/100ML	A213523 002	Mar 09, 2021
		<u>POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER</u>		
AP	BAXTER HLTHCARE	5GM/100ML;150MG/100ML;450MG/100ML	N018008 007	Apr 28, 1982
AP	+! ICU MEDICAL INC	5GM/100ML;149MG/100ML;450MG/100ML	N018362 010	Jul 05, 1983
		<u>POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9%</u>		
AP	FRESENIUS KABI USA	5GM/100ML;149MG/100ML;900MG/100ML	A213445 001	Mar 09, 2021
		<u>POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER</u>		
AP	BAXTER HLTHCARE	5GM/100ML;150MG/100ML;900MG/100ML	N019308 005	Apr 05, 1985
AP	+! ICU MEDICAL INC	5GM/100ML;149MG/100ML;900MG/100ML	N019691 005	Mar 24, 1988

PRESCRIPTION DRUG PRODUCT LIST

DEXTROSE; POTASSIUM CHLORIDE; SODIUM CHLORIDE

INJECTABLE; INJECTION

POTASSIUM CHLORIDE 30MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45%

AP FRESENIUS KABI USA 5GM/100ML;224MG/100ML;450MG/100ML **A213523 003** Mar 09, 2021

POTASSIUM CHLORIDE 30MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER

AP BAXTER HLTHCARE 5GM/100ML;224MG/100ML;450MG/100ML **N018008 008** Apr 28, 1982

AP +! ICU MEDICAL INC 5GM/100ML;224MG/100ML;450MG/100ML **N018362 002**

POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45%

AP FRESENIUS KABI USA 5GM/100ML;298MG/100ML;450MG/100ML **A213523 004** Mar 09, 2021

POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER

AP BAXTER HLTHCARE 5GM/100ML;300MG/100ML;450MG/100ML **N018008 009** Apr 28, 1982

AP +! ICU MEDICAL INC 5GM/100ML;298MG/100ML;450MG/100ML **N018362 003**

POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9%

AP FRESENIUS KABI USA 5GM/100ML;298MG/100ML;900MG/100ML **A213445 002** Mar 09, 2021

POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

AP BAXTER HLTHCARE 5GM/100ML;300MG/100ML;900MG/100ML **N019308 007** Apr 05, 1985

AP +! ICU MEDICAL INC 5GM/100ML;298MG/100ML;900MG/100ML **N019691 009** Mar 24, 1988

POTASSIUM CHLORIDE 0.037% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER

B BRAUN 10GM/100ML;37MG/100ML;200MG/100ML N019630 031 Feb 17, 1988

POTASSIUM CHLORIDE 0.037% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER

B BRAUN 10GM/100ML;37MG/100ML;450MG/100ML N019630 037 Feb 17, 1988

POTASSIUM CHLORIDE 0.037% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

B BRAUN 10GM/100ML;37MG/100ML;900MG/100ML N019630 043 Feb 17, 1988

POTASSIUM CHLORIDE 0.037% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER

B BRAUN 5GM/100ML;37MG/100ML;110MG/100ML N019630 001 Feb 17, 1988

POTASSIUM CHLORIDE 0.037% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER

B BRAUN 5GM/100ML;37MG/100ML;200MG/100ML N019630 007 Feb 17, 1988

POTASSIUM CHLORIDE 0.037% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER

B BRAUN 5GM/100ML;37MG/100ML;330MG/100ML N019630 013 Feb 17, 1988

POTASSIUM CHLORIDE 0.037% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER

B BRAUN 5GM/100ML;37MG/100ML;450MG/100ML N019630 019 Feb 17, 1988

POTASSIUM CHLORIDE 0.037% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

B BRAUN 5GM/100ML;37MG/100ML;900MG/100ML N019630 025 Feb 17, 1988

POTASSIUM CHLORIDE 0.075% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER

B BRAUN 10GM/100ML;75MG/100ML;200MG/100ML N019630 032 Feb 17, 1988

POTASSIUM CHLORIDE 0.075% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER

B BRAUN 10GM/100ML;75MG/100ML;450MG/100ML N019630 038 Feb 17, 1988

POTASSIUM CHLORIDE 0.075% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

B BRAUN 10GM/100ML;75MG/100ML;900MG/100ML N019630 044 Feb 17, 1988

POTASSIUM CHLORIDE 0.075% IN DEXTROSE 3.3% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER

B BRAUN 3.3GM/100ML;75MG/100ML;300MG/100ML N019630 049 May 07, 1992

POTASSIUM CHLORIDE 0.075% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER

B BRAUN 5GM/100ML;75MG/100ML;110MG/100ML N019630 002 Feb 17, 1988

POTASSIUM CHLORIDE 0.11% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER

B BRAUN 10GM/100ML;110MG/100ML;200MG/100ML N019630 033 Feb 17, 1988

POTASSIUM CHLORIDE 0.11% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER

B BRAUN 10GM/100ML;110MG/100ML;450MG/100ML N019630 039 Feb 17, 1988

POTASSIUM CHLORIDE 0.11% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

B BRAUN 10GM/100ML;110MG/100ML;900MG/100ML N019630 045 Feb 17, 1988

POTASSIUM CHLORIDE 0.11% IN DEXTROSE 3.3% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER

B BRAUN 3.3GM/100ML;110MG/100ML;300MG/100ML N019630 050 May 07, 1992

POTASSIUM CHLORIDE 0.11% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER

B BRAUN 5GM/100ML;110MG/100ML;110MG/100ML N019630 003 Feb 17, 1988

POTASSIUM CHLORIDE 0.11% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER

B BRAUN 5GM/100ML;110MG/100ML;200MG/100ML N019630 009 Feb 17, 1988

POTASSIUM CHLORIDE 0.11% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER

B BRAUN 5GM/100ML;110MG/100ML;330MG/100ML N019630 015 Feb 17, 1988

POTASSIUM CHLORIDE 0.11% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER

B BRAUN 5GM/100ML;110MG/100ML;450MG/100ML N019630 021 Feb 17, 1988

POTASSIUM CHLORIDE 0.11% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

B BRAUN 5GM/100ML;110MG/100ML;900MG/100ML N019630 027 Feb 17, 1988

POTASSIUM CHLORIDE 0.15% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER

B BRAUN 10GM/100ML;150MG/100ML;200MG/100ML N019630 034 Feb 17, 1988

POTASSIUM CHLORIDE 0.15% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER

B BRAUN 10GM/100ML;150MG/100ML;450MG/100ML N019630 040 Feb 17, 1988

POTASSIUM CHLORIDE 0.15% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

B BRAUN 10GM/100ML;150MG/100ML;900MG/100ML N019630 046 Feb 17, 1988

POTASSIUM CHLORIDE 0.15% IN DEXTROSE 3.3% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER

B BRAUN 3.3GM/100ML;150MG/100ML;300MG/100ML N019630 051 May 07, 1992

POTASSIUM CHLORIDE 0.15% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER

B BRAUN 5GM/100ML;150MG/100ML;110MG/100ML N019630 004 Feb 17, 1988

PRESCRIPTION DRUG PRODUCT LIST

DEXTROSE; POTASSIUM CHLORIDE; SODIUM CHLORIDE

INJECTABLE; INJECTION

POTASSIUM CHLORIDE 0.22% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER			
B BRAUN	10GM/100ML;220MG/100ML;200MG/100ML	N019630 035	Feb 17, 1988
POTASSIUM CHLORIDE 0.22% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER			
B BRAUN	10GM/100ML;220MG/100ML;450MG/100ML	N019630 041	Feb 17, 1988
POTASSIUM CHLORIDE 0.22% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER			
B BRAUN	10GM/100ML;220MG/100ML;900MG/100ML	N019630 047	Feb 17, 1988
POTASSIUM CHLORIDE 0.22% IN DEXTROSE 3.3% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER			
B BRAUN	3.3GM/100ML;220MG/100ML;300MG/100ML	N019630 052	May 07, 1992
POTASSIUM CHLORIDE 0.22% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER			
B BRAUN	5GM/100ML;220MG/100ML;110MG/100ML	N019630 005	Feb 17, 1988
POTASSIUM CHLORIDE 0.22% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER			
B BRAUN	5GM/100ML;220MG/100ML;200MG/100ML	N019630 011	Feb 17, 1988
POTASSIUM CHLORIDE 0.22% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER			
B BRAUN	5GM/100ML;220MG/100ML;330MG/100ML	N019630 017	Feb 17, 1988
POTASSIUM CHLORIDE 0.22% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER			
B BRAUN	5GM/100ML;220MG/100ML;450MG/100ML	N019630 023	Feb 17, 1988
POTASSIUM CHLORIDE 0.22% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER			
B BRAUN	5GM/100ML;220MG/100ML;900MG/100ML	N019630 029	Feb 17, 1988
POTASSIUM CHLORIDE 0.3% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER			
B BRAUN	10GM/100ML;300MG/100ML;200MG/100ML	N019630 036	Feb 17, 1988
POTASSIUM CHLORIDE 0.3% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER			
B BRAUN	10GM/100ML;300MG/100ML;450MG/100ML	N019630 042	Feb 17, 1988
POTASSIUM CHLORIDE 0.3% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER			
B BRAUN	10GM/100ML;300MG/100ML;900MG/100ML	N019630 048	Feb 17, 1988
POTASSIUM CHLORIDE 0.3% IN DEXTROSE 3.3% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER			
B BRAUN	3.3GM/100ML;300MG/100ML;300MG/100ML	N019630 053	May 07, 1992
POTASSIUM CHLORIDE 0.3% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER			
B BRAUN	5GM/100ML;300MG/100ML;110MG/100ML	N019630 006	Feb 17, 1988
POTASSIUM CHLORIDE 10MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER			
BAXTER HLTHCARE	5GM/100ML;75MG/100ML;900MG/100ML	N019308 004	Apr 05, 1985
	5GM/100ML;150MG/100ML;900MG/100ML	N019308 002	Apr 05, 1985
POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER			
BAXTER HLTHCARE	5GM/100ML;300MG/100ML;900MG/100ML	N019308 003	Apr 05, 1985
POTASSIUM CHLORIDE 30MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER			
BAXTER HLTHCARE	5GM/100ML;224MG/100ML;900MG/100ML	N019308 006	Apr 05, 1985
POTASSIUM CHLORIDE 5MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER			
BAXTER HLTHCARE	5GM/100ML;150MG/100ML;450MG/100ML	N018008 004	
POTASSIUM CHLORIDE 5MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER			
BAXTER HLTHCARE	5GM/100ML;150MG/100ML;900MG/100ML	N019308 001	Apr 05, 1985

DEXTROSE; SODIUM CHLORIDE

INJECTABLE; INJECTION

<u>DEXTROSE 2.5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER</u>			
AP	B BRAUN	<u>2.5GM/100ML;450MG/100ML</u>	<u>N019631 004</u> Feb 24, 1988
AP	+	BAXTER HLTHCARE	<u>2.5GM/100ML;450MG/100ML</u> <u>N016697 001</u>
AP	FRESENIUS KABI USA	<u>2.5GM/100ML;450MG/100ML</u>	<u>A211190 001</u> Dec 20, 2019
<u>DEXTROSE 5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER</u>			
AP	B BRAUN	<u>5GM/100ML;200MG/100ML</u>	<u>N019631 007</u> Feb 24, 1988
<u>DEXTROSE 5% AND SODIUM CHLORIDE 0.225%</u>			
AP	FRESENIUS KABI USA	<u>5GM/100ML;225MG/100ML</u>	<u>A211221 001</u> Sep 15, 2020
<u>DEXTROSE 5% AND SODIUM CHLORIDE 0.225% IN PLASTIC CONTAINER</u>			
AP	+	ICU MEDICAL INC	<u>5GM/100ML;225MG/100ML</u> <u>N017606 001</u>
<u>DEXTROSE 5% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER</u>			
AP	FRESENIUS KABI USA	<u>5GM/100ML;300MG/100ML</u>	<u>A211194 001</u> Aug 26, 2020
AP	+	ICU MEDICAL INC	<u>5GM/100ML;300MG/100ML</u> <u>N017799 001</u>
<u>DEXTROSE 5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER</u>			
AP	B BRAUN	<u>5GM/100ML;330MG/100ML</u>	<u>N019631 008</u> Feb 24, 1988
<u>DEXTROSE 5% AND SODIUM CHLORIDE 0.45%</u>			
AP	FRESENIUS KABI USA	<u>5GM/100ML;450MG/100ML</u>	<u>A211276 001</u> Sep 15, 2020
<u>DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER</u>			
AP	B BRAUN	<u>5GM/100ML;450MG/100ML</u>	<u>N019631 009</u> Feb 24, 1988
AP	+	ICU MEDICAL INC	<u>5GM/100ML;450MG/100ML</u> <u>N017607 001</u>
<u>DEXTROSE 5% AND SODIUM CHLORIDE 0.9%</u>			
AP	FRESENIUS KABI USA	<u>5GM/100ML;900MG/100ML</u>	<u>A211211 001</u> Sep 14, 2020
<u>DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER</u>			
AP	B BRAUN	<u>5GM/100ML;900MG/100ML</u>	<u>N019631 010</u> Feb 24, 1988
AP	+	ICU MEDICAL INC	<u>5GM/100ML;900MG/100ML</u> <u>N017585 001</u>
<u>DEXTROSE 5% IN SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER</u>			
AP	BAXTER HLTHCARE	<u>5GM/100ML;200MG/100ML</u>	<u>N016689 001</u>

PRESCRIPTION DRUG PRODUCT LIST

DEXTROSE; SODIUM CHLORIDE

INJECTABLE; INJECTION

DEXTROSE 5% IN SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER

AP	BAXTER HLTHCARE	<u>5GM/100ML;330MG/100ML</u>	<u>N016687</u>	<u>001</u>	
	<u>DEXTROSE 5% IN SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER</u>				
AP	BAXTER HLTHCARE	<u>5GM/100ML;450MG/100ML</u>	<u>N016683</u>	<u>001</u>	
	<u>DEXTROSE 5% IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER</u>				
AP	BAXTER HLTHCARE	<u>5GM/100ML;900MG/100ML</u>	<u>N016678</u>	<u>001</u>	
	DEXTROSE 10% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER				
	B BRAUN	10GM/100ML;110MG/100ML	N019631	011	Feb 24, 1988
	DEXTROSE 10% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER				
	B BRAUN	10GM/100ML;200MG/100ML	N019631	012	Feb 24, 1988
	DEXTROSE 10% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER				
	B BRAUN	10GM/100ML;330MG/100ML	N019631	013	Feb 24, 1988
	DEXTROSE 10% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER				
	B BRAUN	10GM/100ML;450MG/100ML	N019631	014	Feb 24, 1988
	DEXTROSE 10% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER				
	B BRAUN	10GM/100ML;900MG/100ML	N019631	015	Feb 24, 1988
	DEXTROSE 2.5% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER				
	B BRAUN	2.5GM/100ML;110MG/100ML	N019631	001	Feb 24, 1988
	DEXTROSE 2.5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER				
	B BRAUN	2.5GM/100ML;200MG/100ML	N019631	002	Feb 24, 1988
	DEXTROSE 2.5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER				
	B BRAUN	2.5GM/100ML;330MG/100ML	N019631	003	Feb 24, 1988
	DEXTROSE 2.5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER				
	B BRAUN	2.5GM/100ML;900MG/100ML	N019631	005	Feb 24, 1988
	DEXTROSE 3.3% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER				
	B BRAUN	3.3GM/100ML;300MG/100ML	N019631	016	Jan 19, 1990
	DEXTROSE 5% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER				
	B BRAUN	5GM/100ML;110MG/100ML	N019631	006	Feb 24, 1988

DIATRIZOATE MEGLUMINE

SOLUTION; URETHRAL

CYSTOGRAFIN

	+ BRACCO	30%	N010040	018	
	CYSTOGRAFIN DILUTE				
	+ BRACCO	18%	N010040	022	Nov 09, 1982

DIATRIZOATE MEGLUMINE; DIATRIZOATE SODIUM

SOLUTION; ORAL, RECTAL

DIATRIZOATE MEGLUMINE AND DIATRIZOATE SODIUM

AA	ANDA REPOSITORY	<u>66%;10%</u>	<u>A214201</u>	<u>001</u>	Jun 27, 2022
AA	ANNORA PHARMA	<u>66%;10%</u>	<u>A215049</u>	<u>001</u>	Nov 17, 2023
	<u>GASTROGRAFIN</u>				
AA	+! BRACCO	<u>66%;10%</u>	<u>N011245</u>	<u>003</u>	
	<u>MD-GASTROVIEW</u>				
AA	LIEBEL-FLARSHEIM	<u>66%;10%</u>	<u>A087388</u>	<u>001</u>	

DIAZEPAM

CONCENTRATE; ORAL

DIAZEPAM

AA	CHARTWELL MOLECULAR	<u>5MG/ML</u>	<u>A204433</u>	<u>001</u>	Apr 14, 2014
	<u>DIAZEPAM INTENSOL</u>				
AA	! HIKMA	<u>5MG/ML</u>	<u>A071415</u>	<u>001</u>	Apr 03, 1987
	FILM; BUCCAL				
	LIBERVANT				
	+ AQUESTIVE	5MG	N218623	001	Apr 26, 2024
	+	7.5MG	N218623	002	Apr 26, 2024
	+	10MG	N218623	003	Apr 26, 2024
	+	12.5MG	N218623	004	Apr 26, 2024
	+!	15MG	N218623	005	Apr 26, 2024

GEL; RECTAL

DIASTAT ACUDIAL

AB	+! BAUSCH	<u>10MG/2ML (5MG/ML)</u>	<u>N020648</u>	<u>007</u>	Sep 15, 2005
AB	+!	<u>20MG/4ML (5MG/ML)</u>	<u>N020648</u>	<u>006</u>	Sep 15, 2005

DIAZEPAM

AB	NOVEL LABS INC	<u>10MG/2ML (5MG/ML)</u>	<u>A091076</u>	<u>001</u>	May 30, 2023
AB		<u>20MG/4ML (5MG/ML)</u>	<u>A091076</u>	<u>002</u>	May 30, 2023

DIASTAT

	+! BAUSCH	2.5MG/0.5ML (5MG/ML)	N020648	001	Jul 29, 1997
--	-----------	----------------------	---------	-----	--------------

INJECTABLE; INJECTION

DIAZEPAM

AP	ALEMBOIC	<u>10MG/2ML (5MG/ML)</u>	<u>A218450</u>	<u>001</u>	Apr 25, 2024
AP	! BELOTECA	<u>10MG/2ML (5MG/ML)</u>	<u>A210363</u>	<u>001</u>	Mar 18, 2019

PRESCRIPTION DRUG PRODUCT LIST

DIAZEPAM

INJECTABLE; INJECTION

DIAZEPAM

<u>AP</u>		<u>50MG/10ML (5MG/ML)</u>	<u>A211998 001</u>	Dec 26, 2019
<u>AP</u>	DR REDDYS	<u>10MG/2ML (5MG/ML)</u>	<u>A218422 001</u>	Jul 18, 2024
<u>AP</u>	FRESENIUS KABI USA	<u>10MG/2ML (5MG/ML)</u>	<u>A214745 001</u>	Nov 10, 2022
<u>AP</u>	HIKMA	<u>10MG/2ML (5MG/ML)</u>	<u>A070313 001</u>	Dec 16, 1985
<u>AP</u>		<u>50MG/10ML (5MG/ML)</u>	<u>A070311 001</u>	Dec 16, 1985
<u>AP</u>	HOSPIRA	<u>10MG/2ML (5MG/ML)</u>	<u>A072079 001</u>	Dec 20, 1988
<u>AP</u>	!	<u>50MG/10ML (5MG/ML)</u>	<u>A071583 001</u>	Oct 13, 1987
<u>AP</u>	LONG GROVE PHARMS	<u>10MG/2ML (5MG/ML)</u>	<u>A217178 001</u>	Dec 31, 2024

SOLUTION; ORAL

DIAZEPAM

<u>AA</u>	CHARTWELL MOLECULAR	<u>5MG/5ML</u>	<u>A206477 001</u>	Jun 24, 2016
<u>AA</u>	!	<u>5MG/5ML</u>	<u>A070928 001</u>	Apr 03, 1987

SPRAY; NASAL

VALTOCO

+	NEURELIS INC	5MG/SPRAY	N211635 001	Jan 10, 2020
+		7.5MG/SPRAY	N211635 002	Jan 10, 2020
+	!	10MG/SPRAY	N211635 003	Jan 10, 2020

TABLET; ORAL

DIAZEPAM

<u>AB</u>	AUROBINDO PHARMA LTD	<u>2MG</u>	<u>A217843 001</u>	Dec 14, 2023
<u>AB</u>		<u>5MG</u>	<u>A217843 002</u>	Dec 14, 2023
<u>AB</u>		<u>10MG</u>	<u>A217843 003</u>	Dec 14, 2023
<u>AB</u>	DR REDDYS LABS SA	<u>2MG</u>	<u>A071134 001</u>	Feb 03, 1987
<u>AB</u>		<u>5MG</u>	<u>A071135 001</u>	Feb 03, 1987
<u>AB</u>		<u>10MG</u>	<u>A071136 001</u>	Feb 03, 1987
<u>AB</u>	IVAX SUB TEVA PHARMS	<u>2MG</u>	<u>A071307 001</u>	Dec 10, 1986
<u>AB</u>		<u>5MG</u>	<u>A071321 001</u>	Dec 10, 1986
<u>AB</u>		<u>10MG</u>	<u>A071322 001</u>	Dec 10, 1986
<u>AB</u>	MYLAN	<u>2MG</u>	<u>A070325 002</u>	Sep 04, 1985
<u>AB</u>		<u>5MG</u>	<u>A070325 003</u>	Sep 04, 1985
<u>AB</u>		<u>10MG</u>	<u>A070325 001</u>	Sep 04, 1985
<u>AB</u>	NUVO PHARM	<u>2MG</u>	<u>A070464 002</u>	Sep 10, 2024
<u>AB</u>		<u>5MG</u>	<u>A070464 003</u>	Sep 10, 2024
<u>AB</u>		<u>10MG</u>	<u>A070464 001</u>	Feb 25, 1986
<u>AB</u>	STRIDES PHARMA	<u>2MG</u>	<u>A077749 001</u>	Mar 31, 2006
<u>AB</u>		<u>5MG</u>	<u>A077749 002</u>	Mar 31, 2006
<u>AB</u>		<u>10MG</u>	<u>A077749 003</u>	Mar 31, 2006
<u>VALIUM</u>				
<u>AB</u>	+	WAYLIS THERAP	<u>2MG</u>	<u>N013263 002</u>
<u>AB</u>	+		<u>5MG</u>	<u>N013263 004</u>
<u>AB</u>	+	!	<u>10MG</u>	<u>N013263 006</u>

DIAZOXIDE

SUSPENSION; ORAL

DIAZOXIDE

<u>AB</u>	E5 PHARMA INC	<u>50MG/ML</u>	<u>A211050 001</u>	Dec 20, 2019
<u>AB</u>	NOVITIUM PHARMA	<u>50MG/ML</u>	<u>A210799 001</u>	Jul 08, 2020
<u>PROGLYCEM</u>				
<u>AB</u>	+	!	TEVA BRANDED PHARM	<u>50MG/ML</u>
				<u>N017453 001</u>

DICHLORPHENAMIDE

TABLET; ORAL

KEVEYIS

<u>AB</u>	+	!	XERIS	<u>50MG</u>	<u>N011366 002</u>	Aug 07, 2015
-----------	---	---	-------	-------------	--------------------	--------------

ORMALVI

<u>AB</u>	TORRENT	<u>50MG</u>	<u>A215924 001</u>	Dec 29, 2022
-----------	---------	-------------	--------------------	--------------

DICLOFENAC EPOLAMINE

SYSTEM; TOPICAL

FLECTOR

+	!	IBSA	1.3%	N021234 001	Jan 31, 2007
LICART					
+	!	IBSA INST BIO	1.3%	N206976 001	Dec 19, 2018

PRESCRIPTION DRUG PRODUCT LIST

DICLOFENAC POTASSIUM

CAPSULE;ORAL

DICLOFENAC POTASSIUM

AB	AUROBINDO PHARMA LTD	25MG	A213875 001	Oct 19, 2021
AB	BIONPHARMA	25MG	A204648 001	Feb 23, 2016
AB	STRIDES SOFTGELS	25MG	A210078 001	Dec 03, 2019

ZIPSOR

AB	+! ASSERTIO FOR SOLUTION;ORAL	25MG	N022202 001	Jun 16, 2009
-----------	-------------------------------	-------------	--------------------	--------------

CAMBIA

AB	+! ASSERTIO	50MG	N022165 001	Jun 17, 2009
-----------	-------------	-------------	--------------------	--------------

DICLOFENAC POTASSIUM

AB	ALKEM LABS LTD	50MG	A216635 001	Jul 20, 2022
AB	ANNORA PHARMA	50MG	A215375 001	Mar 04, 2022
AB	ENDO OPERATIONS	50MG	A202964 001	May 02, 2016
AB	TARO	50MG	A218011 001	Jun 14, 2024

TABLET;ORAL

CATAFLAM

AB	AMICI PHARMA	25MG	A076561 002	Jul 21, 2021
AB		50MG	A076561 001	Mar 18, 2004

DICLOFENAC POTASSIUM

AB	NOVAST LABS	50MG	A215585 001	Oct 08, 2021
AB	! RK PHARMA	50MG	A075463 001	Jul 26, 1999
AB	RUBICON	25MG	A075229 002	Sep 16, 2021
AB		50MG	A075229 001	Nov 20, 1998
AB	SENORES PHARMS	25MG	A215787 002	Nov 09, 2023
AB		50MG	A215787 001	Mar 15, 2023
AB	TEVA	50MG	A075219 001	Aug 06, 1998
AB	UMEDICA	50MG	A215750 001	May 11, 2022

DICLOFENAC SODIUM

GEL;TOPICAL

DICLOFENAC SODIUM

AB	ACTAVIS MID ATLANTIC	3%	A206493 001	Dec 02, 2015
AB	ALEMBIC	3%	A212351 001	Jul 27, 2022
AB	AMNEAL	3%	A200936 001	Oct 28, 2013
AB	GLENMARK PHARMS LTD	3%	A208301 001	Sep 13, 2016
AB	PADAGIS ISRAEL	3%	A210893 001	Jul 27, 2018
AB	! TARO	3%	A206298 001	Apr 28, 2016

SOLUTION;TOPICAL

DICLOFENAC SODIUM

AB	ALEMBIC	2%	A212506 001	Nov 29, 2022
AB	AMNEAL	2%	A208198 001	Aug 18, 2022
AB	APOTEX	2%	A207714 001	May 06, 2022
AB	AUROLIFE PHARMA LLC	2%	A213040 001	Feb 03, 2023
AB	! LUPIN PHARMS	2%	A208021 001	Sep 20, 2022
AB	TARO	2%	A208098 001	Sep 29, 2022
AT	AMNEAL PHARMS	1.5%	A206116 001	Sep 02, 2016
AT	EPIC PHARMA LLC	1.5%	A206655 001	Jan 28, 2021
AT	LUPIN LTD	1.5%	A204132 001	Aug 20, 2015
AT	NOVEL LABS INC	1.5%	A205878 001	Dec 09, 2015
AT	! TARO	1.5%	A203818 001	Nov 26, 2014
AT	WATSON LABS INC	1.5%	A202852 001	Nov 24, 2014
AT	ZYDUS LIFESCIENCES	1.5%	A206411 001	Apr 17, 2018

SOLUTION/DROPS;OPHTHALMIC

DICLOFENAC SODIUM

AT	ALTAIRE PHARMS INC	0.1%	A203383 001	Nov 16, 2015
AT	! BAUSCH AND LOMB	0.1%	A078792 001	Dec 28, 2007
AT	RISING	0.1%	A078553 001	Dec 28, 2007
AT	RUBICON	0.1%	A077600 001	Nov 13, 2008
AT	SANDOZ	0.1%	A078031 001	Feb 06, 2008

TABLET, DELAYED RELEASE;ORAL

DICLOFENAC SODIUM

AB	ACTAVIS ELIZABETH	50MG	A074514 001	Mar 26, 1996
AB		75MG	A074514 002	Mar 26, 1996
AB	CARLSBAD	25MG	A075185 002	Nov 13, 1998
AB		50MG	A075185 003	Nov 13, 1998
AB		75MG	A075185 001	Nov 13, 1998
AB	RUBICON	25MG	A216548 001	May 11, 2023
AB		50MG	A216548 002	May 11, 2023
AB		75MG	A216548 003	May 11, 2023

PRESCRIPTION DRUG PRODUCT LIST

DICLOFENAC SODIUM

TABLET, DELAYED RELEASE;ORAL

DICLOFENAC SODIUM

<u>AB</u>	!	UNIQUE	<u>25MG</u>	<u>A090066</u>	<u>001</u>	Dec 01, 2010
<u>AB</u>	!		<u>50MG</u>	<u>A090066</u>	<u>002</u>	Dec 01, 2010
<u>AB</u>	!		<u>75MG</u>	<u>A077863</u>	<u>003</u>	Jun 08, 2007

TABLET, EXTENDED RELEASE;ORAL

DICLOFENAC SODIUM

<u>AB</u>	!	DEXCEL LTD	<u>100MG</u>	<u>A076201</u>	<u>001</u>	Nov 06, 2002
<u>AB</u>		RICONPHARMA LLC	<u>100MG</u>	<u>A216275</u>	<u>001</u>	Sep 23, 2022
<u>AB</u>		VPNA	<u>100MG</u>	<u>A075492</u>	<u>001</u>	Feb 11, 2000

DICLOFENAC SODIUM; MISOPROSTOL

TABLET, DELAYED RELEASE;ORAL

ARTHROTEC

<u>AB</u>	+	PFIZER	<u>50MG;0.2MG</u>	<u>N020607</u>	<u>001</u>	Dec 24, 1997
<u>AB</u>	+	!	<u>75MG;0.2MG</u>	<u>N020607</u>	<u>002</u>	Dec 24, 1997

DICLOFENAC SODIUM AND MISOPROSTOL

<u>AB</u>		ACTAVIS LABS FL INC	<u>50MG;0.2MG</u>	<u>A201089</u>	<u>001</u>	Jul 09, 2012
<u>AB</u>			<u>75MG;0.2MG</u>	<u>A201089</u>	<u>002</u>	Jul 09, 2012
<u>AB</u>		AMNEAL PHARMS	<u>50MG;0.2MG</u>	<u>A203995</u>	<u>001</u>	Nov 25, 2016
<u>AB</u>			<u>75MG;0.2MG</u>	<u>A203995</u>	<u>002</u>	Nov 25, 2016
<u>AB</u>		MICRO LABS	<u>50MG;0.2MG</u>	<u>A204355</u>	<u>001</u>	Jul 15, 2021
<u>AB</u>			<u>75MG;0.2MG</u>	<u>A204355</u>	<u>002</u>	Jul 15, 2021
<u>AB</u>		YUNG SHIN PHARM	<u>50MG;0.2MG</u>	<u>A205143</u>	<u>001</u>	Feb 19, 2020
<u>AB</u>			<u>75MG;0.2MG</u>	<u>A205143</u>	<u>002</u>	Feb 19, 2020

DICLOXACILLIN SODIUM

CAPSULE;ORAL

DICLOXACILLIN SODIUM

<u>AB</u>		AUROBINDO PHARMA	<u>EQ 250MG BASE</u>	<u>A216845</u>	<u>001</u>	Sep 23, 2022
<u>AB</u>			<u>EQ 500MG BASE</u>	<u>A216845</u>	<u>002</u>	Sep 23, 2022
<u>AB</u>		TEVA	<u>EQ 250MG BASE</u>	<u>A062286</u>	<u>001</u>	Jun 03, 1982
<u>AB</u>	!		<u>EQ 500MG BASE</u>	<u>A062286</u>	<u>002</u>	Jun 03, 1982

DICYCLOMINE HYDROCHLORIDE

CAPSULE;ORAL

DICYCLOMINE HYDROCHLORIDE

<u>AB</u>		ANNORA PHARMA	<u>10MG</u>	<u>A218018</u>	<u>001</u>	Jul 01, 2024
<u>AB</u>		AUROBINDO PHARMA USA	<u>10MG</u>	<u>A040319</u>	<u>001</u>	Sep 07, 1999
<u>AB</u>		COREPHARMA	<u>10MG</u>	<u>A216639</u>	<u>001</u>	Mar 24, 2023
<u>AB</u>	!	LANNETT	<u>10MG</u>	<u>A084285</u>	<u>001</u>	
<u>AB</u>		PRINSTON INC	<u>10MG</u>	<u>A217531</u>	<u>001</u>	Aug 30, 2023
<u>AB</u>		TWI PHARMS	<u>10MG</u>	<u>A217054</u>	<u>001</u>	Dec 27, 2022
<u>AB</u>		WATSON LABS	<u>10MG</u>	<u>A085082</u>	<u>001</u>	Jun 19, 1986
<u>AB</u>		WEST WARD	<u>10MG</u>	<u>A040204</u>	<u>001</u>	Feb 28, 1997

INJECTABLE; INJECTION

DICYCLOMINE HYDROCHLORIDE

<u>AP</u>		AM REGENT	<u>10MG/ML</u>	<u>A208353</u>	<u>001</u>	Feb 17, 2017
<u>AP</u>	!	FOSUN PHARMA	<u>10MG/ML</u>	<u>A210979</u>	<u>001</u>	Jul 02, 2018
<u>AP</u>		FRESENIUS KABI USA	<u>10MG/ML</u>	<u>A210257</u>	<u>001</u>	Jan 25, 2019
<u>AP</u>		NEXUS	<u>10MG/ML</u>	<u>A206468</u>	<u>001</u>	Feb 01, 2019
<u>AP</u>		SLATE RUN PHARMA	<u>10MG/ML</u>	<u>A207076</u>	<u>001</u>	Nov 02, 2018
<u>AP</u>		SOMERSET THERAPS LLC	<u>10MG/ML</u>	<u>A214332</u>	<u>001</u>	Nov 01, 2024
<u>AP</u>			<u>10MG/ML</u>	<u>A214333</u>	<u>001</u>	Nov 01, 2024

DICYCLOMINE HYDROCHLORIDE (PRESERVATIVE FREE)

<u>AP</u>	!	HIKMA	<u>10MG/ML</u>	<u>A040465</u>	<u>001</u>	Jun 30, 2003
-----------	---	-------	----------------	----------------	------------	--------------

SYRUP;ORAL

DICYCLOMINE HYDROCHLORIDE

<u>AA</u>	!	ENDO OPERATIONS	<u>10MG/5ML</u>	<u>A040169</u>	<u>001</u>	Mar 24, 2005
<u>AA</u>		HIKMA	<u>10MG/5ML</u>	<u>A212286</u>	<u>001</u>	May 22, 2020
<u>AA</u>		NOVITIUM PHARMA	<u>10MG/5ML</u>	<u>A214721</u>	<u>001</u>	Apr 23, 2021

TABLET;ORAL

DICYCLOMINE HYDROCHLORIDE

<u>AB</u>		ANNORA PHARMA	<u>20MG</u>	<u>A217566</u>	<u>001</u>	May 07, 2024
<u>AB</u>		AUROBINDO PHARMA USA	<u>20MG</u>	<u>A040317</u>	<u>001</u>	Sep 07, 1999
<u>AB</u>		BIONPHARMA	<u>20MG</u>	<u>A217916</u>	<u>001</u>	Aug 04, 2023
<u>AB</u>		COREPHARMA	<u>20MG</u>	<u>A216760</u>	<u>001</u>	Nov 28, 2022
<u>AB</u>		HIKMA PHARMS	<u>20MG</u>	<u>A040161</u>	<u>001</u>	Oct 01, 1996
<u>AB</u>		LANNETT	<u>20MG</u>	<u>A040230</u>	<u>001</u>	Feb 26, 1999
<u>AB</u>		RISING	<u>20MG</u>	<u>A218952</u>	<u>001</u>	Nov 15, 2024

PRESCRIPTION DRUG PRODUCT LIST

DICYCLOMINE HYDROCHLORIDE

TABLET; ORAL

DICYCLOMINE HYDROCHLORIDE

AB	RUBICON	20MG	A216736 001	Dec 14, 2022
AB	TWI PHARMS	20MG	A216782 001	Jun 01, 2023
AB	! WATSON LABS	20MG	A085223 001	Jul 30, 1986

DIENOGEST; ESTRADIOL VALERATE

TABLET; ORAL

NATAZIA

+	BAYER HLTHCARE	N/A, 2MG, 3MG, N/A, N/A; 3MG, 2MG, 2MG, 1MG, N/A	N022252 001	May 06, 2010
---	----------------	--	-------------	--------------

DIETHYLPROPION HYDROCHLORIDE

TABLET; ORAL

DIETHYLPROPION HYDROCHLORIDE

AA	AVANTHI INC	25MG	A201212 001	Dec 22, 2010
AA	! LANNETT CO INC	25MG	A200177 001	Jul 18, 2011
TABLET, EXTENDED RELEASE; ORAL				
DIETHYLPROPION HYDROCHLORIDE				
	! LANNETT CO INC	75MG	A091680 001	Oct 24, 2011

DIFELIKEFALIN ACETATE

SOLUTION; INTRAVENOUS

KORSUVA

+	CARA THERAP	EQ 0.065MG BASE/1.3ML (EQ 0.05MG BASE/ML)	N214916 001	Aug 23, 2021
---	-------------	---	-------------	--------------

DIFLORASONE DIACETATE

CREAM; TOPICAL

DIFLORASONE DIACETATE

BX	! TARO	0.05%	A075508 001	Apr 24, 2000
----	--------	-------	-------------	--------------

OINTMENT; TOPICAL

DIFLORASONE DIACETATE

AB	AVONDALE PHARMS	0.05%	A075374 001	Apr 27, 1999
AB	RISING	0.05%	A207440 001	Feb 27, 2017
AB	! TARO	0.05%	A075331 001	May 14, 1999
AB	THE J MOLNER	0.05%	A210753 001	Jun 12, 2018

DIFLUNISAL

TABLET; ORAL

DIFLUNISAL

AB	HERITAGE PHARMA	500MG	A202845 001	Mar 08, 2012
AB	! TEVA	500MG	A073673 001	Jul 31, 1992
AB	ZYDUS PHARMS	500MG	A203547 001	Jun 16, 2017
	HERITAGE PHARMA	250MG	A202845 002	Aug 16, 2024
		375MG	A202845 003	Aug 16, 2024

DIFLUPREDNATE

EMULSION; OPHTHALMIC

DIFLUPREDNATE

AB	AMNEAL	0.05%	A211526 001	Nov 17, 2021
AB	CAPLIN	0.05%	A219441 001	Dec 13, 2024
AB	CIPLA	0.05%	A211776 001	Aug 09, 2021
AB	DR REDDYS	0.05%	A214894 001	Nov 16, 2022
AB	EPIC PHARMA LLC	0.05%	A207284 001	May 14, 2024

DUREZOL

AB	! SANDOZ	0.05%	N022212 001	Jun 23, 2008
-----------	----------	--------------	--------------------	--------------

DIGOXIN

ELIXIR; ORAL

DIGOXIN

AA	AMICI PHARMA	0.05MG/ML	A215209 001	Mar 11, 2022
AA	! HIKMA	0.05MG/ML	N021648 001	Aug 26, 2004
AA	VISTAPHARM LLC	0.05MG/ML	A213000 001	Oct 04, 2019

INJECTABLE; INJECTION

DIGOXIN

AP	HIKMA	0.25MG/ML	A083391 001	
AP	SANDOZ	0.25MG/ML	A040481 001	Aug 21, 2003

LANOXIN

AP	! COVIS	0.25MG/ML	N009330 002	
LANOXIN PEDIATRIC				
	! COVIS	0.1MG/ML	N009330 004	

TABLET; ORAL

DIGOXIN

AB	AUROBINDO PHARMA LTD	0.0625MG	A214982 001	Feb 08, 2022
-----------	----------------------	-----------------	--------------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

DIGOXIN

TABLET; ORAL

DIGOXIN

<u>AB</u>		<u>0.125MG</u>	<u>A214982</u>	<u>002</u>	Feb 08, 2022
<u>AB</u>		<u>0.25MG</u>	<u>A214982</u>	<u>003</u>	Feb 08, 2022
<u>AB</u>	HIKMA INTL PHARMS	<u>0.125MG</u>	<u>A077002</u>	<u>002</u>	Oct 30, 2007
<u>AB</u>		<u>0.25MG</u>	<u>A077002</u>	<u>001</u>	Oct 30, 2007
<u>AB</u>	NOVITIUM PHARMA	<u>0.0625MG</u>	<u>A215307</u>	<u>003</u>	Aug 25, 2022
<u>AB</u>		<u>0.125MG</u>	<u>A215307</u>	<u>001</u>	Nov 22, 2021
<u>AB</u>		<u>0.25MG</u>	<u>A215307</u>	<u>002</u>	Nov 22, 2021
<u>AB</u>	RISING	<u>0.125MG</u>	<u>A040282</u>	<u>001</u>	Dec 23, 1999
<u>AB</u>		<u>0.25MG</u>	<u>A040282</u>	<u>002</u>	Dec 23, 1999
<u>AB</u>	STEVENS J	<u>0.125MG</u>	<u>A076268</u>	<u>001</u>	Jul 26, 2002
<u>AB</u>		<u>0.25MG</u>	<u>A076268</u>	<u>002</u>	Jul 26, 2002
<u>AB</u>	SUN PHARM INDS INC	<u>0.125MG</u>	<u>A076363</u>	<u>001</u>	Jan 31, 2003
<u>AB</u>		<u>0.25MG</u>	<u>A076363</u>	<u>002</u>	Jan 31, 2003

LANOXIN

<u>AB</u>	+	ADVANZ PHARMA	<u>0.0625MG</u>	<u>N020405</u>	<u>001</u>	Sep 30, 1997
<u>AB</u>	+		<u>0.125MG</u>	<u>N020405</u>	<u>002</u>	Sep 30, 1997
<u>AB</u>	+	!	<u>0.25MG</u>	<u>N020405</u>	<u>004</u>	Sep 30, 1997

DIHYDROERGOTAMINE MESYLATE

INJECTABLE; INJECTION

DIHYDROERGOTAMINE MESYLATE

<u>AP</u>		BAXTER HLTHCARE CORP	<u>1MG/ML</u>	<u>A216747</u>	<u>001</u>	Jan 07, 2025
<u>AP</u>		CIPLA	<u>1MG/ML</u>	<u>A212334</u>	<u>001</u>	Sep 20, 2024
<u>AP</u>		GLAND PHARMA LTD	<u>1MG/ML</u>	<u>A215623</u>	<u>001</u>	Aug 18, 2023
<u>AP</u>		HIKMA	<u>1MG/ML</u>	<u>A040453</u>	<u>001</u>	Jun 09, 2003
<u>AP</u>	!	HIKMA PHARMS	<u>1MG/ML</u>	<u>A206621</u>	<u>001</u>	Sep 15, 2017
<u>AP</u>		PADAGIS US	<u>1MG/ML</u>	<u>A040475</u>	<u>001</u>	Apr 28, 2003
<u>AP</u>		PROVEPHARM SAS	<u>1MG/ML</u>	<u>A212046</u>	<u>001</u>	Jan 07, 2020
<u>AP</u>		SAGENT PHARMS INC	<u>1MG/ML</u>	<u>A207264</u>	<u>001</u>	Jul 11, 2018

SPRAY, METERED; NASAL

DIHYDROERGOTAMINE MESYLATE

<u>AB</u>		CIPLA	<u>0.5MG/SPRAY</u>	<u>A212907</u>	<u>001</u>	May 20, 2020
<u>AB</u>		HIKMA	<u>0.5MG/SPRAY</u>	<u>A211393</u>	<u>001</u>	Feb 28, 2020
<u>AB</u>		RUBICON	<u>0.5MG/SPRAY</u>	<u>A216881</u>	<u>001</u>	Jun 22, 2023

MIGRANAL

<u>AB</u>	+	!	BAUSCH TRUDHESA	<u>0.5MG/SPRAY</u>	<u>N020148</u>	<u>001</u>	Dec 08, 1997
	+	!	IMPTEL PHARMS	0.725MG/SPRAY	N213436	001	Sep 02, 2021

DILTIAZEM HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

DILTIAZEM HYDROCHLORIDE

<u>AB1</u>		DR REDDYS LABS SA	<u>60MG</u>	<u>A215775</u>	<u>001</u>	May 05, 2022
<u>AB1</u>			<u>90MG</u>	<u>A215775</u>	<u>002</u>	May 05, 2022
<u>AB1</u>			<u>120MG</u>	<u>A215775</u>	<u>003</u>	May 05, 2022
<u>AB1</u>		GLENMARK PHARMS LTD	<u>60MG</u>	<u>A212317</u>	<u>001</u>	Mar 22, 2021
<u>AB1</u>			<u>90MG</u>	<u>A212317</u>	<u>002</u>	Mar 22, 2021
<u>AB1</u>			<u>120MG</u>	<u>A212317</u>	<u>003</u>	Mar 22, 2021
<u>AB1</u>		MYLAN	<u>60MG</u>	<u>A074910</u>	<u>001</u>	May 02, 1997
<u>AB1</u>			<u>90MG</u>	<u>A074910</u>	<u>002</u>	May 02, 1997
<u>AB1</u>	!		<u>120MG</u>	<u>A074910</u>	<u>003</u>	May 02, 1997
<u>AB1</u>		TWI PHARMS	<u>60MG</u>	<u>A217377</u>	<u>001</u>	Mar 01, 2023
<u>AB1</u>			<u>90MG</u>	<u>A217377</u>	<u>002</u>	Mar 01, 2023
<u>AB1</u>			<u>120MG</u>	<u>A217377</u>	<u>003</u>	Mar 01, 2023
<u>AB2</u>		ACCORD HLTHCARE	<u>120MG</u>	<u>A206997</u>	<u>001</u>	Apr 28, 2020
<u>AB2</u>			<u>180MG</u>	<u>A206997</u>	<u>002</u>	Apr 28, 2020
<u>AB2</u>			<u>240MG</u>	<u>A206997</u>	<u>003</u>	Apr 28, 2020
<u>AB2</u>		ALEMBIC	<u>120MG</u>	<u>A218587</u>	<u>001</u>	Oct 18, 2024
<u>AB2</u>			<u>180MG</u>	<u>A218587</u>	<u>002</u>	Oct 18, 2024
<u>AB2</u>			<u>240MG</u>	<u>A218587</u>	<u>003</u>	Oct 18, 2024
<u>AB2</u>		APOTEX	<u>120MG</u>	<u>A074943</u>	<u>003</u>	Dec 19, 2000
<u>AB2</u>			<u>180MG</u>	<u>A074943</u>	<u>002</u>	Dec 19, 2000
<u>AB2</u>	!		<u>240MG</u>	<u>A074943</u>	<u>001</u>	Aug 06, 1998
<u>AB2</u>		UTOPIC PHARMS	<u>120MG</u>	<u>A216304</u>	<u>001</u>	Aug 08, 2022
<u>AB2</u>			<u>180MG</u>	<u>A216304</u>	<u>002</u>	Aug 08, 2022
<u>AB2</u>			<u>240MG</u>	<u>A216304</u>	<u>003</u>	Aug 08, 2022

CARDIZEM CD

<u>AB3</u>	+	BAUSCH	<u>120MG</u>	<u>N020062</u>	<u>001</u>	Aug 10, 1992
<u>AB3</u>	+		<u>180MG</u>	<u>N020062</u>	<u>002</u>	Dec 27, 1991
<u>AB3</u>	+		<u>240MG</u>	<u>N020062</u>	<u>003</u>	Dec 27, 1991

PRESCRIPTION DRUG PRODUCT LIST

DILTIAZEM HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

CARDIZEM CD

<u>AB3</u>	+		<u>300MG</u>	<u>N020062</u>	<u>004</u>	Dec 27, 1991
<u>AB3</u>	+	!	<u>360MG</u>	<u>N020062</u>	<u>005</u>	Aug 24, 1999

CARTIA XT

<u>AB3</u>		ACTAVIS LABS FL INC	<u>120MG</u>	<u>A074752</u>	<u>002</u>	Jul 09, 1998
<u>AB3</u>			<u>180MG</u>	<u>A074752</u>	<u>001</u>	Jul 09, 1998
<u>AB3</u>			<u>240MG</u>	<u>A074752</u>	<u>003</u>	Jul 09, 1998
<u>AB3</u>			<u>300MG</u>	<u>A074752</u>	<u>004</u>	Jul 09, 1998

DILTIAZEM HYDROCHLORIDE

<u>AB3</u>		ACTAVIS ELIZABETH	<u>360MG</u>	<u>A202463</u>	<u>001</u>	Dec 07, 2012
<u>AB3</u>		ALEMBIC	<u>120MG</u>	<u>A216968</u>	<u>001</u>	Nov 08, 2024
<u>AB3</u>			<u>180MG</u>	<u>A216968</u>	<u>002</u>	Nov 08, 2024
<u>AB3</u>			<u>240MG</u>	<u>A216968</u>	<u>003</u>	Nov 08, 2024
<u>AB3</u>			<u>300MG</u>	<u>A216968</u>	<u>004</u>	Nov 08, 2024
<u>AB3</u>			<u>360MG</u>	<u>A216968</u>	<u>005</u>	Nov 08, 2024
<u>AB3</u>		ENDO OPERATIONS	<u>120MG</u>	<u>A074984</u>	<u>001</u>	Dec 20, 1999
<u>AB3</u>			<u>180MG</u>	<u>A074984</u>	<u>002</u>	Dec 20, 1999
<u>AB3</u>			<u>240MG</u>	<u>A074984</u>	<u>003</u>	Dec 20, 1999
<u>AB3</u>			<u>300MG</u>	<u>A074984</u>	<u>004</u>	Dec 20, 1999
<u>AB3</u>		NOVAST LABS	<u>120MG</u>	<u>A208783</u>	<u>001</u>	Jun 14, 2019
<u>AB3</u>			<u>180MG</u>	<u>A208783</u>	<u>002</u>	Jun 14, 2019
<u>AB3</u>			<u>240MG</u>	<u>A208783</u>	<u>003</u>	Jun 14, 2019
<u>AB3</u>			<u>300MG</u>	<u>A208783</u>	<u>004</u>	Jun 14, 2019
<u>AB3</u>			<u>360MG</u>	<u>A208783</u>	<u>005</u>	Jun 14, 2019
<u>AB3</u>		SUN PHARM	<u>120MG</u>	<u>A203023</u>	<u>001</u>	Jun 08, 2017
<u>AB3</u>			<u>180MG</u>	<u>A203023</u>	<u>002</u>	Jun 08, 2017
<u>AB3</u>			<u>240MG</u>	<u>A203023</u>	<u>003</u>	Jun 08, 2017
<u>AB3</u>			<u>300MG</u>	<u>A203023</u>	<u>004</u>	Jun 08, 2017
<u>AB3</u>			<u>360MG</u>	<u>A203023</u>	<u>005</u>	Jun 08, 2017
<u>AB3</u>		TWI PHARMS	<u>120MG</u>	<u>A205231</u>	<u>001</u>	Aug 30, 2018
<u>AB3</u>			<u>180MG</u>	<u>A205231</u>	<u>002</u>	Aug 30, 2018
<u>AB3</u>			<u>240MG</u>	<u>A205231</u>	<u>003</u>	Aug 30, 2018
<u>AB3</u>			<u>300MG</u>	<u>A205231</u>	<u>004</u>	Aug 30, 2018
<u>AB3</u>			<u>360MG</u>	<u>A205231</u>	<u>005</u>	Aug 30, 2018
<u>AB3</u>		VALEANT PHARMS NORTH	<u>120MG</u>	<u>A075116</u>	<u>001</u>	Dec 23, 1999
<u>AB3</u>			<u>180MG</u>	<u>A075116</u>	<u>002</u>	Dec 23, 1999
<u>AB3</u>			<u>240MG</u>	<u>A075116</u>	<u>003</u>	Dec 23, 1999
<u>AB3</u>			<u>300MG</u>	<u>A075116</u>	<u>004</u>	Dec 23, 1999
<u>AB3</u>		ZYDUS PHARMS	<u>120MG</u>	<u>A206534</u>	<u>001</u>	Aug 08, 2017
<u>AB3</u>			<u>180MG</u>	<u>A206534</u>	<u>002</u>	Aug 08, 2017
<u>AB3</u>			<u>240MG</u>	<u>A206534</u>	<u>003</u>	Aug 08, 2017
<u>AB3</u>			<u>300MG</u>	<u>A206534</u>	<u>004</u>	Aug 08, 2017
<u>AB3</u>			<u>360MG</u>	<u>A206534</u>	<u>005</u>	Aug 08, 2017
<u>AB4</u>		CHARTWELL RX	<u>120MG</u>	<u>A091022</u>	<u>001</u>	Sep 28, 2012
<u>AB4</u>			<u>180MG</u>	<u>A091022</u>	<u>002</u>	Sep 28, 2012
<u>AB4</u>			<u>240MG</u>	<u>A091022</u>	<u>003</u>	Sep 28, 2012
<u>AB4</u>			<u>300MG</u>	<u>A091022</u>	<u>004</u>	Sep 28, 2012
<u>AB4</u>			<u>360MG</u>	<u>A091022</u>	<u>005</u>	Sep 28, 2012
<u>AB4</u>			<u>420MG</u>	<u>A091022</u>	<u>006</u>	Sep 28, 2012
<u>AB4</u>		ZYDUS PHARMS	<u>120MG</u>	<u>A206641</u>	<u>001</u>	Aug 11, 2017
<u>AB4</u>			<u>180MG</u>	<u>A206641</u>	<u>002</u>	Aug 11, 2017
<u>AB4</u>			<u>240MG</u>	<u>A206641</u>	<u>003</u>	Aug 11, 2017
<u>AB4</u>			<u>300MG</u>	<u>A206641</u>	<u>004</u>	Aug 11, 2017
<u>AB4</u>			<u>360MG</u>	<u>A206641</u>	<u>005</u>	Aug 11, 2017
<u>AB4</u>			<u>420MG</u>	<u>A206641</u>	<u>006</u>	Aug 11, 2017

TIAZAC

<u>AB4</u>	+	BAUSCH	<u>120MG</u>	<u>N020401</u>	<u>001</u>	Sep 11, 1995
<u>AB4</u>	+		<u>180MG</u>	<u>N020401</u>	<u>002</u>	Sep 11, 1995
<u>AB4</u>	+		<u>240MG</u>	<u>N020401</u>	<u>003</u>	Sep 11, 1995
<u>AB4</u>	+		<u>300MG</u>	<u>N020401</u>	<u>004</u>	Sep 11, 1995
<u>AB4</u>	+		<u>360MG</u>	<u>N020401</u>	<u>005</u>	Sep 11, 1995
<u>AB4</u>	+	!	<u>420MG</u>	<u>N020401</u>	<u>006</u>	Oct 16, 1998

INJECTABLE; INJECTION

DILTIAZEM HYDROCHLORIDE

<u>AP</u>		EUGIA PHARMA	<u>5MG/ML</u>	<u>A216552</u>	<u>001</u>	Mar 29, 2023
<u>AP</u>		HIKMA	<u>5MG/ML</u>	<u>A078538</u>	<u>001</u>	Dec 17, 2008
<u>AP</u>		HOSPIRA	<u>5MG/ML</u>	<u>A074941</u>	<u>001</u>	Apr 15, 1998
<u>AP</u>		RISING	<u>5MG/ML</u>	<u>A075086</u>	<u>001</u>	Apr 09, 1998
<u>AP</u>	!	SAGENT	<u>5MG/ML</u>	<u>A074617</u>	<u>001</u>	Feb 28, 1996
	!	HOSPIRA	100MG/VIAL	A075853	001	Dec 17, 2002

PRESCRIPTION DRUG PRODUCT LIST

DILTIAZEM HYDROCHLORIDE

TABLET; ORAL

CARDIZEM

<u>AB</u>	+	BAUSCH	<u>30MG</u>	<u>N018602</u>	<u>001</u>	Nov 05, 1982
<u>AB</u>	+		<u>60MG</u>	<u>N018602</u>	<u>002</u>	Nov 05, 1982
<u>AB</u>	+		<u>90MG</u>	<u>N018602</u>	<u>003</u>	Dec 08, 1986
<u>AB</u>	+	!	<u>120MG</u>	<u>N018602</u>	<u>004</u>	Dec 08, 1986

DILTIAZEM HYDROCHLORIDE

<u>AB</u>		SCIEGEN PHARMS INC	<u>30MG</u>	<u>A216521</u>	<u>001</u>	Sep 23, 2022
<u>AB</u>			<u>60MG</u>	<u>A216521</u>	<u>002</u>	Sep 23, 2022
<u>AB</u>			<u>90MG</u>	<u>A216521</u>	<u>003</u>	Sep 23, 2022
<u>AB</u>			<u>120MG</u>	<u>A216521</u>	<u>004</u>	Sep 23, 2022
<u>AB</u>		TEVA	<u>30MG</u>	<u>A074185</u>	<u>001</u>	May 31, 1995
<u>AB</u>			<u>60MG</u>	<u>A074185</u>	<u>002</u>	May 31, 1995
<u>AB</u>			<u>90MG</u>	<u>A074185</u>	<u>003</u>	May 31, 1995
<u>AB</u>			<u>120MG</u>	<u>A074185</u>	<u>004</u>	May 31, 1995

TABLET, EXTENDED RELEASE; ORAL

CARDIZEM LA

<u>AB</u>	+	BAUSCH	<u>120MG</u>	<u>N021392</u>	<u>001</u>	Feb 06, 2003
<u>AB</u>	+		<u>180MG</u>	<u>N021392</u>	<u>002</u>	Feb 06, 2003
<u>AB</u>	+		<u>240MG</u>	<u>N021392</u>	<u>003</u>	Feb 06, 2003
<u>AB</u>	+		<u>300MG</u>	<u>N021392</u>	<u>004</u>	Feb 06, 2003
<u>AB</u>	+		<u>360MG</u>	<u>N021392</u>	<u>005</u>	Feb 06, 2003
<u>AB</u>	+	!	<u>420MG</u>	<u>N021392</u>	<u>006</u>	Feb 06, 2003

DILTIAZEM HYDROCHLORIDE

<u>AB</u>		ACTAVIS LABS FL INC	<u>120MG</u>	<u>A077686</u>	<u>006</u>	Mar 15, 2010
<u>AB</u>			<u>180MG</u>	<u>A077686</u>	<u>005</u>	Mar 15, 2010
<u>AB</u>			<u>240MG</u>	<u>A077686</u>	<u>004</u>	Mar 15, 2010
<u>AB</u>			<u>300MG</u>	<u>A077686</u>	<u>003</u>	Mar 15, 2010
<u>AB</u>			<u>360MG</u>	<u>A077686</u>	<u>002</u>	Mar 15, 2010
<u>AB</u>			<u>420MG</u>	<u>A077686</u>	<u>001</u>	Mar 15, 2010
<u>AB</u>		AMTA	<u>120MG</u>	<u>A216439</u>	<u>001</u>	Mar 07, 2023
<u>AB</u>			<u>180MG</u>	<u>A216439</u>	<u>002</u>	Mar 07, 2023
<u>AB</u>			<u>240MG</u>	<u>A216439</u>	<u>003</u>	Mar 07, 2023
<u>AB</u>			<u>300MG</u>	<u>A216439</u>	<u>004</u>	Mar 07, 2023
<u>AB</u>			<u>360MG</u>	<u>A216439</u>	<u>005</u>	Mar 07, 2023
<u>AB</u>			<u>420MG</u>	<u>A216439</u>	<u>006</u>	Mar 07, 2023
<u>AB</u>		SCIEGEN PHARMS INC	<u>120MG</u>	<u>A216327</u>	<u>001</u>	Apr 06, 2023
<u>AB</u>			<u>180MG</u>	<u>A216327</u>	<u>002</u>	Apr 06, 2023
<u>AB</u>			<u>240MG</u>	<u>A216327</u>	<u>003</u>	Apr 06, 2023
<u>AB</u>			<u>300MG</u>	<u>A216327</u>	<u>004</u>	Apr 06, 2023
<u>AB</u>			<u>360MG</u>	<u>A216327</u>	<u>005</u>	Apr 06, 2023
<u>AB</u>			<u>420MG</u>	<u>A216327</u>	<u>006</u>	Apr 06, 2023

DIMENHYDRINATE

INJECTABLE; INJECTION

DIMENHYDRINATE

!	FRESENIUS KABI USA	50MG/ML	A040519	001	Jun 23, 2004
---	--------------------	---------	---------	-----	--------------

DIMERCAPROL

INJECTABLE; INJECTION

BAL

+	!	PROVEPHARM SAS	10%	N005939	001
---	---	----------------	-----	---------	-----

DIMETHYL FUMARATE

CAPSULE, DELAYED RELEASE; ORAL

DIMETHYL FUMARATE

<u>AB</u>		ACCORD HLTHCARE	<u>120MG</u>	<u>A210499</u>	<u>001</u>	Sep 24, 2020
<u>AB</u>			<u>240MG</u>	<u>A210499</u>	<u>002</u>	Sep 24, 2020
<u>AB</u>		ALKEM LABS LTD	<u>120MG</u>	<u>A210440</u>	<u>001</u>	Sep 24, 2020
<u>AB</u>			<u>240MG</u>	<u>A210440</u>	<u>002</u>	Sep 24, 2020
<u>AB</u>		AMNEAL	<u>120MG</u>	<u>A210402</u>	<u>001</u>	Sep 24, 2020
<u>AB</u>			<u>240MG</u>	<u>A210402</u>	<u>002</u>	Sep 24, 2020
<u>AB</u>		AUROBINDO PHARMA	<u>120MG</u>	<u>A210385</u>	<u>001</u>	Dec 22, 2022
<u>AB</u>			<u>240MG</u>	<u>A210385</u>	<u>002</u>	Dec 22, 2022
<u>AB</u>		CIPLA	<u>120MG</u>	<u>A210305</u>	<u>001</u>	Sep 24, 2020
<u>AB</u>			<u>240MG</u>	<u>A210305</u>	<u>002</u>	Sep 24, 2020
<u>AB</u>		GLENMARK PHARMS LTD	<u>120MG</u>	<u>A210309</u>	<u>001</u>	Oct 06, 2020
<u>AB</u>			<u>240MG</u>	<u>A210309</u>	<u>002</u>	Oct 06, 2020
<u>AB</u>		HETERO LABS LTD III	<u>120MG</u>	<u>A210500</u>	<u>001</u>	Sep 24, 2020
<u>AB</u>			<u>240MG</u>	<u>A210500</u>	<u>002</u>	Sep 24, 2020
<u>AB</u>		LUPIN	<u>120MG</u>	<u>A210226</u>	<u>001</u>	Oct 05, 2020
<u>AB</u>			<u>240MG</u>	<u>A210226</u>	<u>002</u>	Oct 05, 2020
<u>AB</u>		MACLEODS PHARMS LTD	<u>120MG</u>	<u>A210377</u>	<u>001</u>	Jun 26, 2024

PRESCRIPTION DRUG PRODUCT LIST

DIMETHYL FUMARATE

CAPSULE, DELAYED RELEASE;ORAL

DIMETHYL FUMARATE

<u>AB</u>		<u>240MG</u>	<u>A210377</u>	<u>002</u>	Jun 26, 2024
<u>AB</u>	MSN	<u>120MG</u>	<u>A210460</u>	<u>001</u>	Sep 24, 2020
<u>AB</u>		<u>240MG</u>	<u>A210460</u>	<u>002</u>	Sep 24, 2020
<u>AB</u>	MYLAN	<u>120MG</u>	<u>A210531</u>	<u>001</u>	Aug 17, 2020
<u>AB</u>		<u>240MG</u>	<u>A210531</u>	<u>002</u>	Aug 17, 2020
<u>AB</u>	PRINSTON INC	<u>120MG</u>	<u>A210414</u>	<u>001</u>	Oct 18, 2022
<u>AB</u>		<u>240MG</u>	<u>A210414</u>	<u>002</u>	Oct 18, 2022
<u>AB</u>	SOLA PHARMS	<u>120MG</u>	<u>A210436</u>	<u>001</u>	Mar 26, 2021
<u>AB</u>		<u>240MG</u>	<u>A210436</u>	<u>002</u>	Mar 26, 2021
<u>AB</u>	TORRENT	<u>120MG</u>	<u>A210390</u>	<u>001</u>	Jan 06, 2025
<u>AB</u>		<u>240MG</u>	<u>A210390</u>	<u>002</u>	Jan 06, 2025
<u>AB</u>	TWI PHARMS	<u>120MG</u>	<u>A210382</u>	<u>001</u>	Oct 14, 2020
<u>AB</u>		<u>240MG</u>	<u>A210382</u>	<u>002</u>	Oct 14, 2020

TECFIDERA

<u>AB</u>	+	BIOGEN INC	<u>120MG</u>	<u>N204063</u>	<u>001</u>	Mar 27, 2013
<u>AB</u>	+	!	<u>240MG</u>	<u>N204063</u>	<u>002</u>	Mar 27, 2013

DIMETHYL SULFOXIDE

SOLUTION;INTRAVESICAL

RIMSO-50

+! MYLAN INSTITUTIONAL 50% N017788 001

DINOPROSTONE

GEL;ENDOCERVICAL

PREPIDIL

+! PFIZER 0.5MG/3GM N019617 001 Dec 09, 1992

INSERT, EXTENDED RELEASE;VAGINAL

CERVIDIL

+! FERRING PHARMS INC 10MG N020411 001 Mar 30, 1995

DIPHENHYDRAMINE HYDROCHLORIDE

ELIXIR;ORAL

DIPHENHYDRAMINE HYDROCHLORIDE

+! PHARM ASSOC 12.5MG/5ML A087513 001 Feb 10, 1982

INJECTABLE;INJECTION

DIPHENHYDRAMINE HYDROCHLORIDE

<u>AP</u>		APP PHARMS	<u>50MG/ML</u>	<u>A040466</u>	<u>001</u>	May 28, 2002
<u>AP</u>		GLAND PHARMA LTD	<u>50MG/ML</u>	<u>A218448</u>	<u>001</u>	Mar 20, 2024
<u>AP</u>		MICRO LABS	<u>50MG/ML</u>	<u>A205723</u>	<u>001</u>	Aug 22, 2018
<u>AP</u>	!	MYLAN INSTITUTIONAL	<u>50MG/ML</u>	<u>A040498</u>	<u>001</u>	Jul 12, 2005
<u>AP</u>	+	WEST-WARD PHARMS	<u>50MG/ML</u>	<u>A080817</u>	<u>002</u>	
		INT				

DIPHENHYDRAMINE HYDROCHLORIDE PRESERVATIVE FREE

<u>AP</u>		FRESENIUS KABI USA	<u>50MG/ML</u>	<u>A091526</u>	<u>001</u>	Mar 26, 2013
-----------	--	--------------------	----------------	----------------	------------	--------------

DIPYRIDAMOLE

INJECTABLE;INJECTION

DIPYRIDAMOLE

<u>AP</u>	!	CHARTWELL	<u>5MG/ML</u>	<u>A074939</u>	<u>001</u>	Apr 13, 1998
		INJECTABLE				
<u>AP</u>		HIKMA	<u>5MG/ML</u>	<u>A074521</u>	<u>001</u>	Oct 18, 1996

TABLET;ORAL

DIPYRIDAMOLE

<u>AB</u>		BARR	<u>25MG</u>	<u>A087184</u>	<u>001</u>	Oct 03, 1990
<u>AB</u>			<u>50MG</u>	<u>A087716</u>	<u>001</u>	Oct 03, 1990
<u>AB</u>			<u>75MG</u>	<u>A087717</u>	<u>001</u>	Oct 03, 1990
<u>AB</u>		IMPAX LABS	<u>25MG</u>	<u>A040782</u>	<u>001</u>	Jul 18, 2007
<u>AB</u>			<u>50MG</u>	<u>A040782</u>	<u>002</u>	Jul 18, 2007
<u>AB</u>	!		<u>75MG</u>	<u>A040782</u>	<u>003</u>	Jul 18, 2007
<u>AB</u>		OXFORD PHARMS	<u>25MG</u>	<u>A040542</u>	<u>001</u>	Apr 21, 2006
<u>AB</u>			<u>50MG</u>	<u>A040542</u>	<u>002</u>	Apr 21, 2006
<u>AB</u>			<u>75MG</u>	<u>A040542</u>	<u>003</u>	Apr 21, 2006
<u>AB</u>		RISING	<u>25MG</u>	<u>A040733</u>	<u>001</u>	Feb 13, 2007
<u>AB</u>			<u>50MG</u>	<u>A040733</u>	<u>002</u>	Feb 13, 2007
<u>AB</u>			<u>75MG</u>	<u>A040733</u>	<u>003</u>	Feb 13, 2007
<u>AB</u>		ZYDUS PHARMS USA	<u>25MG</u>	<u>A040874</u>	<u>001</u>	Jan 28, 2008
		INC				
<u>AB</u>			<u>50MG</u>	<u>A040874</u>	<u>002</u>	Jan 28, 2008
<u>AB</u>			<u>75MG</u>	<u>A040874</u>	<u>003</u>	Jan 28, 2008

PRESCRIPTION DRUG PRODUCT LIST

DIROXIMEL FUMARATE

CAPSULE, DELAYED RELEASE;ORAL

VUMERITY

+! BIOGEN INC

231MG

N211855 001 Oct 29, 2019

DISOPYRAMIDE PHOSPHATE

CAPSULE;ORAL

DISOPYRAMIDE PHOSPHATEAB DR REDDYS LABS SAEQ 100MG BASEA070173 001 May 31, 1985ABEQ 150MG BASEA070173 002 May 31, 1985AB TEVAEQ 100MG BASEA070101 001 Feb 22, 1985ABEQ 150MG BASEA070102 001 Feb 22, 1985NORPACEAB + PFIZEREQ 100MG BASEN017447 001AB +!EQ 150MG BASEN017447 002

CAPSULE, EXTENDED RELEASE;ORAL

NORPACE CR

+ PFIZER

EQ 100MG BASE

N018655 001 Jul 20, 1982

+!

EQ 150MG BASE

N018655 002 Jul 20, 1982

DISULFIRAM

TABLET;ORAL

DISULFIRAMAB ALVOGEN250MGA091681 001 Aug 08, 2013AB CHARTWELL MOLECULES250MGA091563 001 Dec 31, 2012AB !500MGA091563 002 Dec 31, 2012AB SIGMAPHARM LABS LLC250MGA091619 001 Mar 28, 2011AB500MGA091619 002 Mar 28, 2011DIVALPROEX SODIUM

CAPSULE, DELAYED REL PELLETS;ORAL

DEPAKOTEAB +! ABBVIEEQ 125MG VALPROIC ACIDN019680 001 Sep 12, 1989DIVALPROEX SODIUMAB AJANTA PHARMA LTDEQ 125MG VALPROIC ACIDA213181 001 Mar 02, 2020AB ALEMBICEQ 125MG VALPROIC ACIDA218793 001 Dec 19, 2024AB DR REDDYS LABS LTDEQ 125MG VALPROIC ACIDA078979 001 Jan 23, 2009AB ZYDUS PHARMS USAEQ 125MG VALPROIC ACIDA078919 001 Jan 27, 2009

INC

TABLET, DELAYED RELEASE;ORAL

DEPAKOTEAB + ABBVIEEQ 125MG VALPROIC ACIDN018723 003 Oct 26, 1984AB +EQ 250MG VALPROIC ACIDN018723 001 Mar 10, 1983AB +!EQ 500MG VALPROIC ACIDN018723 002 Mar 10, 1983DIVALPROEX SODIUMAB APOTEXEQ 125MG VALPROIC ACIDA077615 003 Jul 29, 2008ABEQ 250MG VALPROIC ACIDA077615 002 Jul 29, 2008ABEQ 500MG VALPROIC ACIDA077615 001 Jul 29, 2008AB AUROBINDO PHARMAEQ 125MG VALPROIC ACIDA090554 001 Apr 21, 2011

LTD

ABEQ 250MG VALPROIC ACIDA090554 002 Apr 21, 2011ABEQ 500MG VALPROIC ACIDA090554 003 Apr 21, 2011AB CHARTWELL RXEQ 125MG VALPROIC ACIDA077296 001 Jul 31, 2008ABEQ 250MG VALPROIC ACIDA077296 002 Jul 31, 2008ABEQ 500MG VALPROIC ACIDA077296 003 Jul 31, 2008AB DR REDDYS LABS LTDEQ 125MG VALPROIC ACIDA078755 001 Jul 29, 2008ABEQ 250MG VALPROIC ACIDA078755 002 Jul 29, 2008ABEQ 500MG VALPROIC ACIDA078755 003 Jul 29, 2008AB INVATECHEQ 125MG VALPROIC ACIDA078290 003 Jul 29, 2008ABEQ 250MG VALPROIC ACIDA078290 002 Jul 29, 2008ABEQ 500MG VALPROIC ACIDA078290 001 Jul 29, 2008AB LUPINEQ 125MG VALPROIC ACIDA078790 001 Jul 29, 2008ABEQ 250MG VALPROIC ACIDA078790 002 Jul 29, 2008ABEQ 500MG VALPROIC ACIDA078790 003 Jul 29, 2008AB ORBION PHARMSEQ 125MG VALPROIC ACIDA078853 001 Nov 25, 2008ABEQ 250MG VALPROIC ACIDA078853 002 Nov 25, 2008ABEQ 500MG VALPROIC ACIDA078853 003 Nov 25, 2008AB PRINSTON INCEQ 125MG VALPROIC ACIDA090210 001 Nov 30, 2009ABEQ 250MG VALPROIC ACIDA090210 002 Nov 30, 2009ABEQ 500MG VALPROIC ACIDA090210 003 Nov 30, 2009AB SUN PHARM INDSEQ 125MG VALPROIC ACIDA078597 001 Jul 29, 2008ABEQ 250MG VALPROIC ACIDA078597 002 Jul 29, 2008ABEQ 500MG VALPROIC ACIDA078597 003 Jul 29, 2008AB UNICHEM LABS LTDEQ 125MG VALPROIC ACIDA079163 001 Apr 05, 2011

PRESCRIPTION DRUG PRODUCT LIST

DIVALPROEX SODIUM

TABLET, DELAYED RELEASE;ORAL

DIVALPROEX SODIUM

<u>AB</u>		<u>EQ 250MG VALPROIC ACID</u>	<u>A079163 002</u>	Apr 05, 2011
<u>AB</u>		<u>EQ 500MG VALPROIC ACID</u>	<u>A079163 003</u>	Apr 05, 2011
<u>AB</u>	UPSHER SMITH LABS	<u>EQ 125MG VALPROIC ACID</u>	<u>A078182 001</u>	Jul 29, 2008
<u>AB</u>		<u>EQ 250MG VALPROIC ACID</u>	<u>A078182 002</u>	Jul 29, 2008
<u>AB</u>		<u>EQ 500MG VALPROIC ACID</u>	<u>A078182 003</u>	Jul 29, 2008
<u>AB</u>	ZYDUS PHARMS USA INC	<u>EQ 125MG VALPROIC ACID</u>	<u>A077100 001</u>	Mar 05, 2009
<u>AB</u>		<u>EQ 250MG VALPROIC ACID</u>	<u>A077100 002</u>	Mar 05, 2009
<u>AB</u>		<u>EQ 500MG VALPROIC ACID</u>	<u>A077100 003</u>	Mar 05, 2009

TABLET, EXTENDED RELEASE;ORAL

DEPAKOTE ER

<u>AB</u>	+	ABBVIE	<u>EQ 250MG VALPROIC ACID</u>	<u>N021168 002</u>	May 31, 2002
<u>AB</u>	+		<u>EQ 500MG VALPROIC ACID</u>	<u>N021168 001</u>	Aug 04, 2000

DIVALPROEX SODIUM

<u>AB</u>		AMNEAL PHARMS	<u>EQ 250MG VALPROIC ACID</u>	<u>A203730 001</u>	May 29, 2015
<u>AB</u>	!		<u>EQ 500MG VALPROIC ACID</u>	<u>A203730 002</u>	May 29, 2015
<u>AB</u>		ANNORA PHARMA	<u>EQ 250MG VALPROIC ACID</u>	<u>A215527 001</u>	Sep 26, 2023
<u>AB</u>			<u>EQ 500MG VALPROIC ACID</u>	<u>A215527 002</u>	Sep 26, 2023
<u>AB</u>		AUROBINDO PHARMA LTD	<u>EQ 250MG VALPROIC ACID</u>	<u>A202419 001</u>	Jun 02, 2014
<u>AB</u>			<u>EQ 500MG VALPROIC ACID</u>	<u>A202419 002</u>	Jun 02, 2014
<u>AB</u>		DR REDDYS LABS LTD	<u>EQ 250MG VALPROIC ACID</u>	<u>A090161 001</u>	Mar 15, 2012
<u>AB</u>		LUPIN LTD	<u>EQ 250MG VALPROIC ACID</u>	<u>A209286 001</u>	Oct 18, 2019
<u>AB</u>			<u>EQ 500MG VALPROIC ACID</u>	<u>A209286 002</u>	Oct 18, 2019
<u>AB</u>		MYLAN	<u>EQ 250MG VALPROIC ACID</u>	<u>A077567 001</u>	Jan 29, 2009
<u>AB</u>			<u>EQ 500MG VALPROIC ACID</u>	<u>A077567 002</u>	Jan 29, 2009
<u>AB</u>		REDDYS	<u>EQ 500MG VALPROIC ACID</u>	<u>A090070 001</u>	Mar 12, 2012
<u>AB</u>		UNICHEM	<u>EQ 250MG VALPROIC ACID</u>	<u>A214643 001</u>	Feb 25, 2022
<u>AB</u>			<u>EQ 500MG VALPROIC ACID</u>	<u>A214643 002</u>	Feb 25, 2022
<u>AB</u>		WOCKHARDT	<u>EQ 250MG VALPROIC ACID</u>	<u>A078705 002</u>	Feb 10, 2009
<u>AB</u>			<u>EQ 500MG VALPROIC ACID</u>	<u>A078705 001</u>	Aug 04, 2009
<u>AB</u>		ZYDUS PHARMS USA INC	<u>EQ 250MG VALPROIC ACID</u>	<u>A078239 001</u>	Feb 27, 2009
<u>AB</u>			<u>EQ 500MG VALPROIC ACID</u>	<u>A078239 002</u>	Aug 04, 2009

DOBUTAMINE HYDROCHLORIDE

INJECTABLE; INJECTION

DOBUTAMINE HYDROCHLORIDE

<u>AP</u>		HIKMA	<u>EQ 12.5MG BASE/ML</u>	<u>A074277 001</u>	Oct 31, 1994
<u>AP</u>	!	HOSPIRA	<u>EQ 12.5MG BASE/ML</u>	<u>A074086 001</u>	Nov 29, 1993
<u>AP</u>		SLATE RUN PHARMA	<u>EQ 12.5MG BASE/ML</u>	<u>A216131 001</u>	Dec 21, 2022

DOBUTAMINE HYDROCHLORIDE IN DEXTROSE 5% IN PLASTIC CONTAINER

<u>AP</u>	+	BAXTER HLTHCARE	<u>EQ 50MG BASE/100ML</u>	<u>N020255 001</u>	Oct 19, 1993
<u>AP</u>	+		<u>EQ 100MG BASE/100ML</u>	<u>N020255 003</u>	Oct 19, 1993
<u>AP</u>	+		<u>EQ 200MG BASE/100ML</u>	<u>N020255 004</u>	Oct 19, 1993
<u>AP</u>	+		<u>EQ 400MG BASE/100ML</u>	<u>N020255 005</u>	Oct 19, 1993
<u>AP</u>	+	HOSPIRA	<u>EQ 50MG BASE/100ML</u>	<u>N020201 003</u>	Oct 19, 1993
<u>AP</u>	+		<u>EQ 100MG BASE/100ML</u>	<u>N020201 002</u>	Oct 19, 1993
<u>AP</u>	+		<u>EQ 200MG BASE/100ML</u>	<u>N020201 001</u>	Oct 19, 1993
<u>AP</u>	+		<u>EQ 400MG BASE/100ML</u>	<u>N020201 006</u>	Jul 07, 1994

DOCETAXEL

INJECTABLE; INJECTION

DOCETAXEL

<u>AP</u>	+	ACCORD HLTHCARE	<u>20MG/ML (20MG/ML)</u>	<u>N201195 003</u>	Apr 20, 2012
<u>AP</u>	+		<u>80MG/4ML (20MG/ML)</u>	<u>N201195 004</u>	Apr 20, 2012
<u>AP</u>	+		<u>160MG/8ML (20MG/ML)</u>	<u>N201195 005</u>	Apr 20, 2012
<u>AP</u>		ACTAVIS	<u>20MG/ML (20MG/ML)</u>	<u>N203551 001</u>	Apr 12, 2013
<u>AP</u>			<u>80MG/4ML (20MG/ML)</u>	<u>N203551 002</u>	Apr 12, 2013
<u>AP</u>			<u>160MG/8ML (20MG/ML)</u>	<u>N203551 004</u>	Sep 21, 2015
<u>AP</u>		ALEMBIC	<u>20MG/2ML (10MG/ML)</u>	<u>A215744 001</u>	Feb 28, 2023
<u>AP</u>			<u>80MG/8ML (10MG/ML)</u>	<u>A215744 002</u>	Feb 28, 2023
<u>AP</u>			<u>160MG/16ML (10MG/ML)</u>	<u>A215744 003</u>	Feb 28, 2023
<u>AP</u>		AMNEAL	<u>20MG/ML (20MG/ML)</u>	<u>A209640 001</u>	Jan 19, 2018
<u>AP</u>			<u>80MG/4ML (20MG/ML)</u>	<u>A209640 002</u>	Jan 19, 2018
<u>AP</u>			<u>160MG/8ML (20MG/ML)</u>	<u>A209640 003</u>	Jan 19, 2018
<u>AP</u>		DR REDDYS	<u>20MG/ML (20MG/ML)</u>	<u>A204193 001</u>	Nov 05, 2014
<u>AP</u>			<u>80MG/4ML (20MG/ML)</u>	<u>A204193 002</u>	Nov 05, 2014
<u>AP</u>		EUGIA PHARMA	<u>20MG/2ML (10MG/ML)</u>	<u>A214575 001</u>	Jun 25, 2021
<u>AP</u>			<u>80MG/8ML (10MG/ML)</u>	<u>A214575 002</u>	Jun 25, 2021
<u>AP</u>			<u>160MG/16ML (10MG/ML)</u>	<u>A214575 003</u>	Jun 25, 2021

PRESCRIPTION DRUG PRODUCT LIST

DOCETAXEL

INJECTABLE; INJECTION

DOCETAXEL

<u>AP</u>	GLAND PHARMA LTD	<u>20MG/2ML (10MG/ML)</u>	<u>A213510 001</u>	Jul 01, 2021
<u>AP</u>		<u>80MG/8ML (10MG/ML)</u>	<u>A213510 002</u>	Jul 01, 2021
<u>AP</u>		<u>160MG/16ML (10MG/ML)</u>	<u>A213510 003</u>	Jul 01, 2021
<u>AP</u>	HENGRUI PHARMA	<u>20MG/ML (20MG/ML)</u>	<u>A207252 001</u>	Aug 09, 2017
<u>AP</u>		<u>80MG/4ML (20MG/ML)</u>	<u>A207252 002</u>	Aug 09, 2017
<u>AP</u>		<u>160MG/8ML (20MG/ML)</u>	<u>A207252 003</u>	Aug 09, 2017
<u>AP</u>	HERITAGE	<u>80MG/4ML (20MG/ML)</u>	<u>A216677 002</u>	Feb 28, 2024
<u>AP</u>		<u>160MG/8ML (20MG/ML)</u>	<u>A216677 003</u>	Nov 26, 2024
<u>AP</u>		<u>20MG/ML (20MG/ML)</u>	<u>A216677 001</u>	Feb 28, 2024
<u>AP</u>	HIKMA	<u>20MG/ML (20MG/ML)</u>	<u>A204490 001</u>	Jan 14, 2021
<u>AP</u>		<u>80MG/4ML (20MG/ML)</u>	<u>A204490 002</u>	Jan 14, 2021
<u>AP</u>	+! HOSPIRA INC	<u>20MG/2ML (10MG/ML)</u>	<u>N022234 001</u>	Mar 08, 2011
<u>AP</u>	+!	<u>80MG/8ML (10MG/ML)</u>	<u>N022234 002</u>	Mar 08, 2011
<u>AP</u>	+!	<u>160MG/16ML (10MG/ML)</u>	<u>N022234 003</u>	Mar 08, 2011
<u>AP</u>	MEITHEAL	<u>20MG/2ML (10MG/ML)</u>	<u>A209634 001</u>	Aug 24, 2018
<u>AP</u>		<u>80MG/8ML (10MG/ML)</u>	<u>A209634 002</u>	Aug 24, 2018
<u>AP</u>		<u>160MG/16ML (10MG/ML)</u>	<u>A209634 003</u>	Aug 24, 2018
<u>AP</u>	MYLAN LABS LTD	<u>20MG/2ML (10MG/ML)</u>	<u>A210072 001</u>	Jul 02, 2018
<u>AP</u>		<u>80MG/8ML (10MG/ML)</u>	<u>A210848 001</u>	Jul 06, 2018
<u>AP</u>		<u>160MG/8ML (20MG/ML)</u>	<u>A208137 001</u>	Apr 01, 2019
<u>AP</u>		<u>160MG/16ML (10MG/ML)</u>	<u>A208859 001</u>	Apr 30, 2018
<u>AP</u>	NOVAST LABS	<u>80MG/8ML (10MG/ML)</u>	<u>A207563 002</u>	Aug 31, 2017
<u>AP</u>		<u>160MG/16ML (10MG/ML)</u>	<u>A207563 003</u>	Aug 31, 2017
<u>AP</u>	SANDOZ	<u>20MG/2ML (10MG/ML)</u>	<u>N201525 001</u>	Jun 29, 2011
<u>AP</u>		<u>80MG/8ML (10MG/ML)</u>	<u>N201525 002</u>	Jun 29, 2011
<u>AP</u>		<u>160MG/16ML (10MG/ML)</u>	<u>N201525 003</u>	Jun 29, 2011
<u>AP</u>	SHILPA	<u>20MG/ML (20MG/ML)</u>	<u>A210327 001</u>	May 16, 2019
<u>AP</u>	+!	<u>20MG/ML (20MG/ML)</u>	<u>N205934 001</u>	Dec 22, 2015
<u>AP</u>		<u>80MG/4ML (20MG/ML)</u>	<u>A210327 002</u>	May 16, 2019
<u>AP</u>	+!	<u>80MG/4ML (20MG/ML)</u>	<u>N205934 002</u>	Dec 22, 2015
<u>AP</u>		<u>160MG/8ML (20MG/ML)</u>	<u>A210327 003</u>	May 16, 2019
<u>AP</u>	+!	<u>160MG/8ML (20MG/ML)</u>	<u>N205934 003</u>	Dec 22, 2015

TAXOTERE

<u>AP</u>	+! SANOFI AVENTIS US	<u>20MG/ML (20MG/ML)</u>	<u>N020449 003</u>	Aug 03, 2010
<u>AP</u>	+!	<u>80MG/4ML (20MG/ML)</u>	<u>N020449 004</u>	Aug 02, 2010
<u>AP</u>	+!	<u>160MG/8ML (20MG/ML)</u>	<u>N020449 005</u>	Apr 13, 2012

DOCETAXEL

	ACTAVIS	140MG/7ML (20MG/ML)	N203551 003	Apr 12, 2013
	+ HOSPIRA INC	20MG/ML (20MG/ML)	N022234 004	Jun 23, 2016
	+ HOSPIRA INC	80MG/4ML (20MG/ML)	N022234 005	Jun 23, 2016
	+ HOSPIRA INC	160MG/8ML (20MG/ML)	N022234 007	Jan 24, 2017

SOLUTION; INTRAVENOUS

BEIZRAY

	+! ZHUHAI	80MG/4ML (20MG/ML)	N218711 001	Oct 23, 2024
--	-----------	--------------------	-------------	--------------

DOCIVYX

	+! AVYXA HOLDINGS	20MG/2ML (10MG/ML)	N215813 001	Nov 22, 2022
	+! AVYXA HOLDINGS	80MG/8ML (10MG/ML)	N215813 002	Nov 22, 2022
	+! AVYXA HOLDINGS	160MG/16ML (10MG/ML)	N215813 003	Nov 22, 2022

DOFETILIDE

CAPSULE; ORAL

DOFETILIDE

<u>AB</u>	AUROBINDO PHARMA LTD	<u>0.125MG</u>	<u>A210740 001</u>	Jan 22, 2019
<u>AB</u>		<u>0.25MG</u>	<u>A210740 002</u>	Jan 22, 2019
<u>AB</u>		<u>0.5MG</u>	<u>A210740 003</u>	Jan 22, 2019

DOFETILIDE

<u>AB</u>	ACCORD HLTHCARE	<u>0.125MG</u>	<u>A213338 001</u>	Jun 19, 2020
<u>AB</u>		<u>0.25MG</u>	<u>A213338 002</u>	Jun 19, 2020
<u>AB</u>		<u>0.5MG</u>	<u>A213338 003</u>	Jun 19, 2020
<u>AB</u>	BIONPHARMA	<u>0.125MG</u>	<u>A208625 001</u>	Apr 10, 2018
<u>AB</u>		<u>0.25MG</u>	<u>A208625 002</u>	Apr 10, 2018
<u>AB</u>		<u>0.5MG</u>	<u>A208625 003</u>	Apr 10, 2018
<u>AB</u>	DR REDDYS LABS SA	<u>0.125MG</u>	<u>A207058 001</u>	Jun 06, 2016
<u>AB</u>		<u>0.25MG</u>	<u>A207058 002</u>	Jun 06, 2016
<u>AB</u>		<u>0.5MG</u>	<u>A207058 003</u>	Jun 06, 2016
<u>AB</u>	GRANULES	<u>0.125MG</u>	<u>A212750 001</u>	Oct 14, 2021
<u>AB</u>		<u>0.25MG</u>	<u>A212750 002</u>	Oct 14, 2021
<u>AB</u>		<u>0.5MG</u>	<u>A212750 003</u>	Oct 14, 2021
<u>AB</u>	MSN	<u>0.125MG</u>	<u>A213220 001</u>	Jan 29, 2020

PRESCRIPTION DRUG PRODUCT LIST

DOFETILIDE

CAPSULE; ORAL

DOFETILIDE

<u>AB</u>		<u>0.25MG</u>	<u>A213220</u>	<u>002</u>	Jan 29, 2020
<u>AB</u>		<u>0.5MG</u>	<u>A213220</u>	<u>003</u>	Jan 29, 2020
<u>AB</u>	SIGMAPHARM LABS LLC	<u>0.125MG</u>	<u>A207746</u>	<u>001</u>	Mar 26, 2018
<u>AB</u>		<u>0.25MG</u>	<u>A207746</u>	<u>002</u>	Mar 26, 2018
<u>AB</u>		<u>0.5MG</u>	<u>A207746</u>	<u>003</u>	Mar 26, 2018
<u>AB</u>	STRIDES PHARMA	<u>0.125MG</u>	<u>A208519</u>	<u>001</u>	Oct 09, 2018
<u>AB</u>		<u>0.25MG</u>	<u>A208519</u>	<u>002</u>	Oct 09, 2018
<u>AB</u>		<u>0.5MG</u>	<u>A208519</u>	<u>003</u>	Oct 09, 2018
<u>AB</u>	SUN PHARM	<u>0.125MG</u>	<u>A210466</u>	<u>001</u>	Oct 09, 2018
<u>AB</u>		<u>0.25MG</u>	<u>A210466</u>	<u>002</u>	Oct 09, 2018
<u>AB</u>		<u>0.5MG</u>	<u>A210466</u>	<u>003</u>	Oct 09, 2018

TIKOSYN

<u>AB</u>	+ PFIZER	<u>0.125MG</u>	<u>N020931</u>	<u>001</u>	Oct 01, 1999
<u>AB</u>	+	<u>0.25MG</u>	<u>N020931</u>	<u>002</u>	Oct 01, 1999
<u>AB</u>	+	<u>0.5MG</u>	<u>N020931</u>	<u>003</u>	Oct 01, 1999

DOLUTEGRAVIR SODIUM

TABLET; ORAL

TIVICAY

	+! VIIV HLTHCARE	EQ 50MG BASE	N204790	001	Aug 12, 2013
TABLET, FOR SUSPENSION; ORAL					
<u>TIVICAY PD</u>					
	+! VIIV HLTHCARE	EQ 5MG BASE	N213983	001	Jun 12, 2020

DOLUTEGRAVIR SODIUM; LAMIVUDINE

TABLET; ORAL

DOVATO

	+! VIIV HLTHCARE	EQ 50MG BASE; 300MG	N211994	001	Apr 08, 2019
--	------------------	---------------------	---------	-----	--------------

DOLUTEGRAVIR SODIUM; RILPIVIRINE HYDROCHLORIDE

TABLET; ORAL

JULUCA

	+! VIIV HLTHCARE	EQ 50MG BASE; EQ 25MG BASE	N210192	001	Nov 21, 2017
--	------------------	----------------------------	---------	-----	--------------

DONEPEZIL HYDROCHLORIDE

SYSTEM; TRANSDERMAL

ADLARITY

	+ CORIUM	5MG/DAY	N212304	001	Mar 11, 2022
	+!	10MG/DAY	N212304	002	Mar 11, 2022

TABLET; ORAL

ARICEPT

<u>AB</u>	+ EISAI INC	<u>5MG</u>	<u>N020690</u>	<u>002</u>	Nov 25, 1996
<u>AB</u>	+	<u>10MG</u>	<u>N020690</u>	<u>001</u>	Nov 25, 1996
<u>AB</u>	+	<u>23MG</u>	<u>N022568</u>	<u>001</u>	Jul 23, 2010

DONEPEZIL HYDROCHLORIDE

<u>AB</u>	ALEMBIC PHARMS LTD	<u>5MG</u>	<u>A201724</u>	<u>001</u>	Feb 25, 2013
<u>AB</u>		<u>10MG</u>	<u>A201724</u>	<u>002</u>	Feb 25, 2013
<u>AB</u>	AUROBINDO	<u>5MG</u>	<u>A090056</u>	<u>001</u>	May 31, 2011
<u>AB</u>		<u>10MG</u>	<u>A090056</u>	<u>002</u>	May 31, 2011
<u>AB</u>	CADILA PHARMS LTD	<u>5MG</u>	<u>A204609</u>	<u>001</u>	Sep 19, 2017
<u>AB</u>		<u>10MG</u>	<u>A204609</u>	<u>002</u>	Sep 19, 2017
<u>AB</u>	CHARTWELL RX	<u>5MG</u>	<u>A090425</u>	<u>001</u>	May 31, 2011
<u>AB</u>		<u>10MG</u>	<u>A090425</u>	<u>002</u>	May 31, 2011
<u>AB</u>	CIPLA LTD	<u>5MG</u>	<u>A077518</u>	<u>001</u>	May 31, 2011
<u>AB</u>		<u>10MG</u>	<u>A077518</u>	<u>002</u>	May 31, 2011
<u>AB</u>	DEXCEL	<u>23MG</u>	<u>A203713</u>	<u>001</u>	Feb 19, 2016
<u>AB</u>	DR REDDYS	<u>23MG</u>	<u>A202723</u>	<u>001</u>	Jul 24, 2013
<u>AB</u>	DR REDDYS LABS LTD	<u>5MG</u>	<u>A201001</u>	<u>001</u>	May 31, 2011
<u>AB</u>		<u>10MG</u>	<u>A201001</u>	<u>002</u>	May 31, 2011
<u>AB</u>	ESJAY PHARMA	<u>5MG</u>	<u>A090551</u>	<u>001</u>	May 31, 2011
<u>AB</u>		<u>10MG</u>	<u>A090551</u>	<u>002</u>	May 31, 2011
<u>AB</u>	HETERO LABS LTD V	<u>5MG</u>	<u>A203034</u>	<u>001</u>	Jan 30, 2015
<u>AB</u>		<u>10MG</u>	<u>A203034</u>	<u>002</u>	Jan 30, 2015
<u>AB</u>	INDICUS PHARMA	<u>5MG</u>	<u>A201634</u>	<u>001</u>	Jun 13, 2012
<u>AB</u>		<u>10MG</u>	<u>A201634</u>	<u>002</u>	Jun 13, 2012
<u>AB</u>	JUBILANT GENERICS	<u>5MG</u>	<u>A090768</u>	<u>001</u>	May 31, 2011
<u>AB</u>		<u>10MG</u>	<u>A090768</u>	<u>002</u>	May 31, 2011
<u>AB</u>	LUPIN LTD	<u>23MG</u>	<u>A202782</u>	<u>001</u>	Oct 30, 2015
<u>AB</u>	MACLEODS PHARMS LTD	<u>5MG</u>	<u>A201146</u>	<u>001</u>	Aug 17, 2012
<u>AB</u>		<u>10MG</u>	<u>A201146</u>	<u>002</u>	Aug 17, 2012
<u>AB</u>		<u>23MG</u>	<u>A202631</u>	<u>001</u>	Jan 22, 2014

PRESCRIPTION DRUG PRODUCT LIST

DONEPEZIL HYDROCHLORIDE

TABLET; ORAL

DONEPEZIL HYDROCHLORIDE

<u>AB</u>	PRINSTON INC	<u>5MG</u>	<u>A200292 001</u>	May 31, 2011
<u>AB</u>		<u>10MG</u>	<u>A200292 002</u>	May 31, 2011
<u>AB</u>	RISING	<u>5MG</u>	<u>A202114 001</u>	Jul 05, 2013
<u>AB</u>		<u>10MG</u>	<u>A202114 002</u>	Jul 05, 2013
<u>AB</u>	SCIEGEN PHARMS INC	<u>5MG</u>	<u>A203907 001</u>	Oct 29, 2014
<u>AB</u>		<u>10MG</u>	<u>A203907 002</u>	Oct 29, 2014
<u>AB</u>	STRIDES PHARMA	<u>5MG</u>	<u>A078662 001</u>	May 31, 2011
<u>AB</u>		<u>10MG</u>	<u>A078662 002</u>	May 31, 2011
<u>AB</u>	TORRENT PHARMS	<u>5MG</u>	<u>A090686 001</u>	May 31, 2011
<u>AB</u>		<u>10MG</u>	<u>A090686 002</u>	May 31, 2011
<u>AB</u>	TWI PHARMS	<u>23MG</u>	<u>A203104 001</u>	Oct 29, 2014
<u>AB</u>	ZYDUS LIFESCIENCES	<u>5MG</u>	<u>A090100 001</u>	Oct 24, 2012
<u>AB</u>		<u>10MG</u>	<u>A090100 002</u>	Oct 24, 2012
<u>AB</u>	ZYDUS PHARMS	<u>23MG</u>	<u>A203162 001</u>	Aug 31, 2017

TABLET, ORALLY DISINTEGRATING; ORAL

DONEPEZIL HYDROCHLORIDE

<u>AB</u>	HISUN PHARM HANGZHOU	<u>5MG</u>	<u>A205269 001</u>	Jul 27, 2018
<u>AB</u>		<u>10MG</u>	<u>A205269 002</u>	Jul 27, 2018
<u>AB</u>	MACLEODS PHARMS LTD	<u>5MG</u>	<u>A201787 001</u>	Dec 14, 2012
<u>AB</u>	!	<u>10MG</u>	<u>A201787 002</u>	Dec 14, 2012

DONEPEZIL HYDROCHLORIDE; MEMANTINE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

MEMANTINE HYDROCHLORIDE AND DONEPEZIL HYDROCHLORIDE

<u>AB</u>	AMNEAL PHARMS	<u>10MG; 14MG</u>	<u>A208328 001</u>	Jan 27, 2017
<u>AB</u>		<u>10MG; 28MG</u>	<u>A208328 002</u>	Jan 27, 2017
<u>AB</u>	ANI PHARMS	<u>10MG; 21MG</u>	<u>A208237 001</u>	Dec 15, 2023

NAMZARIC

<u>AB</u>	+ ABBVIE	<u>10MG; 14MG</u>	<u>N206439 001</u>	Dec 23, 2014
<u>AB</u>	+	<u>10MG; 21MG</u>	<u>N206439 004</u>	Jul 18, 2016
<u>AB</u>	+!	<u>10MG; 28MG</u>	<u>N206439 002</u>	Dec 23, 2014
	+	10MG; 7MG	N206439 003	Jul 18, 2016

DOPAMINE HYDROCHLORIDE

INJECTABLE; INJECTION

DOPAMINE HYDROCHLORIDE

<u>AP</u>	HIKMA INTL PHARMS	<u>40MG/ML</u>	<u>A207707 001</u>	Apr 11, 2018
<u>AP</u>		<u>80MG/ML</u>	<u>A207707 002</u>	Apr 11, 2018
<u>AP</u>	+! HOSPIRA	<u>40MG/ML</u>	<u>N018132 001</u>	
<u>AP</u>	+!	<u>80MG/100ML</u>	<u>N018132 002</u>	Feb 04, 1982
<u>AP</u>	+!	<u>80MG/ML</u>	<u>N018132 004</u>	Jul 09, 1982
<u>AP</u>	+!	<u>160MG/100ML</u>	<u>N018132 003</u>	Feb 04, 1982

DOPAMINE HYDROCHLORIDE IN DEXTROSE 5% IN PLASTIC CONTAINER

<u>AP</u>	+! BAXTER HLTHCARE	<u>80MG/100ML</u>	<u>N019615 001</u>	Mar 27, 1987
<u>AP</u>	+!	<u>160MG/100ML</u>	<u>N019615 002</u>	Mar 27, 1987
<u>AP</u>	+!	<u>320MG/100ML</u>	<u>N019615 003</u>	Mar 27, 1987
<u>AP</u>	+! HOSPIRA	<u>80MG/100ML</u>	<u>N018826 001</u>	Sep 30, 1983
<u>AP</u>	+!	<u>160MG/100ML</u>	<u>N018826 002</u>	Sep 30, 1983
<u>AP</u>	+!	<u>320MG/100ML</u>	<u>N018826 003</u>	Sep 30, 1983
	+! BAXTER HLTHCARE	640MG/100ML	N019615 004	Mar 27, 1987

DORAVIRINE

TABLET; ORAL

PIFELTRO

+!	MSD MERCK CO	100MG	N210806 001	Aug 30, 2018
----	--------------	-------	-------------	--------------

DORAVIRINE; LAMIVUDINE; TENOFOVIR DISOPROXIL FUMARATE

TABLET; ORAL

DELSTRIGO

+!	MSD MERCK CO	100MG; 300MG; 300MG	N210807 001	Aug 30, 2018
----	--------------	---------------------	-------------	--------------

DORZOLAMIDE HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

DORZOLAMIDE HYDROCHLORIDE

<u>AT</u>	ALEMBIC	<u>EQ 2% BASE</u>	<u>A212639 001</u>	Aug 09, 2019
<u>AT</u>	BAUSCH AND LOMB	<u>EQ 2% BASE</u>	<u>A090143 001</u>	Jun 25, 2009
<u>AT</u>	FDC LTD	<u>EQ 2% BASE</u>	<u>A205294 001</u>	Jan 24, 2019
<u>AT</u>	GLAND PHARMA LTD	<u>EQ 2% BASE</u>	<u>A215660 001</u>	Jan 27, 2022
<u>AT</u>	INDOCO	<u>EQ 2% BASE</u>	<u>A202053 001</u>	Sep 11, 2014
<u>AT</u>	! MICRO LABS	<u>EQ 2% BASE</u>	<u>A204778 001</u>	Nov 08, 2019
<u>AT</u>	RUBICON	<u>EQ 2% BASE</u>	<u>A078395 001</u>	Oct 28, 2008

PRESCRIPTION DRUG PRODUCT LIST

DORZOLAMIDE HYDROCHLORIDE

SOLUTION/DROPS;OPHTHALMIC

DORZOLAMIDE HYDROCHLORIDE

<u>AT</u>	SANDOZ	<u>EQ 2% BASE</u>	<u>A078748 001</u>	Nov 06, 2008
<u>AT</u>		<u>EQ 2% BASE</u>	<u>A078981 001</u>	Apr 13, 2009

DORZOLAMIDE HYDROCHLORIDE; TIMOLOL MALEATE

SOLUTION/DROPS;OPHTHALMIC

COSOPT

<u>AT1</u>	+	THEA PHARMA	<u>EQ 2% BASE;EQ 0.5% BASE</u>	<u>N020869 001</u>	Apr 07, 1998
------------	---	-------------	--------------------------------	--------------------	--------------

DORZOLAMIDE HYDROCHLORIDE AND TIMOLOL MALEATE

<u>AT1</u>		ALEMBIC	<u>EQ 2% BASE;EQ 0.5% BASE</u>	<u>A213099 001</u>	May 04, 2021
<u>AT1</u>		BAUSCH AND LOMB	<u>EQ 2% BASE;EQ 0.5% BASE</u>	<u>A090037 001</u>	Jul 14, 2009
<u>AT1</u>		EPIC PHARMA LLC	<u>EQ 2% BASE;EQ 0.5% BASE</u>	<u>A203058 001</u>	Sep 22, 2014
<u>AT1</u>		EUGIA PHARMA	<u>EQ 2% BASE;EQ 0.5% BASE</u>	<u>A207629 001</u>	May 14, 2021
<u>AT1</u>		FDC LTD	<u>EQ 2% BASE;EQ 0.5% BASE</u>	<u>A205295 001</u>	Jun 13, 2019
<u>AT1</u>		GLAND PHARMA LTD	<u>EQ 2% BASE;EQ 0.5% BASE</u>	<u>A215520 001</u>	Sep 19, 2022
<u>AT1</u>		INDOCO	<u>EQ 2% BASE;EQ 0.5% BASE</u>	<u>A202054 001</u>	Sep 03, 2014
<u>AT1</u>		MICRO LABS	<u>EQ 2% BASE;EQ 0.5% BASE</u>	<u>A204777 001</u>	May 28, 2020
<u>AT1</u>		SANDOZ	<u>EQ 2% BASE;EQ 0.5% BASE</u>	<u>A078749 001</u>	Nov 06, 2008
<u>AT1</u>			<u>EQ 2% BASE;EQ 0.5% BASE</u>	<u>A090604 001</u>	Nov 18, 2009
<u>AT1</u>		SOMERSET	<u>EQ 2% BASE;EQ 0.5% BASE</u>	<u>A207523 001</u>	Jun 25, 2019

COSOPT PF

<u>AT2</u>	+	THEA PHARMA	<u>EQ 2% BASE;EQ 0.5% BASE</u>	<u>N202667 001</u>	Feb 01, 2012
------------	---	-------------	--------------------------------	--------------------	--------------

DORZOLAMIDE HYDROCHLORIDE AND TIMOLOL MALEATE

<u>AT2</u>		EUGIA PHARMA	<u>EQ 2% BASE;EQ 0.5% BASE</u>	<u>A207630 001</u>	Jul 24, 2018
<u>AT2</u>		INGENUS PHARMS LLC	<u>EQ 2% BASE;EQ 0.5% BASE</u>	<u>A217260 001</u>	May 01, 2023
<u>AT2</u>		MICRO LABS	<u>EQ 2% BASE;EQ 0.5% BASE</u>	<u>A215936 001</u>	Jan 25, 2022

DOXAPRAM HYDROCHLORIDE

INJECTABLE;INJECTION

DOPRAM

<u>AP</u>	+	HIKMA	<u>20MG/ML</u>	<u>N014879 001</u>	
-----------	---	-------	----------------	--------------------	--

DOXAPRAM HYDROCHLORIDE

<u>AP</u>		CHARTWELL INJECTABLE	<u>20MG/ML</u>	<u>A076266 001</u>	Jan 10, 2003
-----------	--	-------------------------	----------------	--------------------	--------------

DOXAZOSIN MESYLATE

TABLET;ORAL

CARDURA

<u>AB</u>	+	VIATRIS	<u>EQ 1MG BASE</u>	<u>N019668 001</u>	Nov 02, 1990
<u>AB</u>	+		<u>EQ 2MG BASE</u>	<u>N019668 002</u>	Nov 02, 1990
<u>AB</u>	+		<u>EQ 4MG BASE</u>	<u>N019668 003</u>	Nov 02, 1990
<u>AB</u>	+		<u>EQ 8MG BASE</u>	<u>N019668 004</u>	Nov 02, 1990

DOXAZOSIN MESYLATE

<u>AB</u>		ACCORD HLTHCARE	<u>EQ 1MG BASE</u>	<u>A202824 001</u>	Jun 11, 2014
<u>AB</u>			<u>EQ 2MG BASE</u>	<u>A202824 002</u>	Jun 11, 2014
<u>AB</u>			<u>EQ 4MG BASE</u>	<u>A202824 003</u>	Jun 11, 2014
<u>AB</u>			<u>EQ 8MG BASE</u>	<u>A202824 004</u>	Jun 11, 2014
<u>AB</u>		APOTEX	<u>EQ 1MG BASE</u>	<u>A075580 001</u>	Oct 18, 2000
<u>AB</u>			<u>EQ 2MG BASE</u>	<u>A075580 002</u>	Oct 18, 2000
<u>AB</u>			<u>EQ 4MG BASE</u>	<u>A075580 003</u>	Oct 18, 2000
<u>AB</u>			<u>EQ 8MG BASE</u>	<u>A075580 004</u>	Oct 18, 2000
<u>AB</u>		AUROBINDO PHARMA USA	<u>EQ 1MG BASE</u>	<u>A075509 001</u>	Oct 19, 2000
<u>AB</u>			<u>EQ 2MG BASE</u>	<u>A075509 002</u>	Oct 19, 2000
<u>AB</u>			<u>EQ 4MG BASE</u>	<u>A075509 003</u>	Oct 19, 2000
<u>AB</u>			<u>EQ 8MG BASE</u>	<u>A075509 004</u>	Oct 19, 2000
<u>AB</u>		HERITAGE PHARMA	<u>EQ 1MG BASE</u>	<u>A205210 001</u>	Feb 13, 2018
<u>AB</u>			<u>EQ 2MG BASE</u>	<u>A205210 002</u>	Feb 13, 2018
<u>AB</u>			<u>EQ 4MG BASE</u>	<u>A205210 003</u>	Feb 13, 2018
<u>AB</u>			<u>EQ 8MG BASE</u>	<u>A205210 004</u>	Feb 13, 2018
<u>AB</u>		RISING	<u>EQ 1MG BASE</u>	<u>A212727 001</u>	Mar 15, 2022
<u>AB</u>			<u>EQ 2MG BASE</u>	<u>A212727 002</u>	Mar 15, 2022
<u>AB</u>			<u>EQ 4MG BASE</u>	<u>A212727 003</u>	Mar 15, 2022
<u>AB</u>			<u>EQ 8MG BASE</u>	<u>A212727 004</u>	Mar 15, 2022
<u>AB</u>		TEVA	<u>EQ 1MG BASE</u>	<u>A075536 001</u>	Oct 18, 2000
<u>AB</u>			<u>EQ 2MG BASE</u>	<u>A075536 002</u>	Oct 18, 2000
<u>AB</u>			<u>EQ 4MG BASE</u>	<u>A075536 003</u>	Oct 18, 2000
<u>AB</u>			<u>EQ 8MG BASE</u>	<u>A075536 004</u>	Oct 18, 2000
<u>AB</u>		UNICHEM	<u>EQ 1MG BASE</u>	<u>A212329 001</u>	Jan 10, 2024
<u>AB</u>			<u>EQ 2MG BASE</u>	<u>A212329 002</u>	Jan 10, 2024
<u>AB</u>			<u>EQ 4MG BASE</u>	<u>A212329 003</u>	Jan 10, 2024
<u>AB</u>			<u>EQ 8MG BASE</u>	<u>A212329 004</u>	Jan 10, 2024

PRESCRIPTION DRUG PRODUCT LIST

DOXAZOSIN MESYLATE

TABLET;ORAL

DOXAZOSIN MESYLATE

<u>AB</u>	UPSHER SMITH LABS	<u>EQ 1MG BASE</u>	<u>A209013 001</u>	Apr 17, 2018
<u>AB</u>		<u>EQ 2MG BASE</u>	<u>A209013 002</u>	Apr 17, 2018
<u>AB</u>		<u>EQ 4MG BASE</u>	<u>A209013 003</u>	Apr 17, 2018
<u>AB</u>		<u>EQ 8MG BASE</u>	<u>A209013 004</u>	Apr 17, 2018
<u>AB</u>	ZYDUS PHARMS	<u>EQ 1MG BASE</u>	<u>A208719 001</u>	Jul 07, 2017
<u>AB</u>		<u>EQ 2MG BASE</u>	<u>A208719 002</u>	Jul 07, 2017
<u>AB</u>		<u>EQ 4MG BASE</u>	<u>A208719 003</u>	Jul 07, 2017
<u>AB</u>		<u>EQ 8MG BASE</u>	<u>A208719 004</u>	Jul 07, 2017

TABLET, EXTENDED RELEASE;ORAL

CARDURA XL

+ VIATRIS

EQ 4MG BASE

N021269 001 Feb 22, 2005

+!

EQ 8MG BASE

N021269 002 Feb 22, 2005

DOXEPIN HYDROCHLORIDE

CAPSULE;ORAL

DOXEPIN HYDROCHLORIDE

<u>AB</u>	ADAPTIS	<u>EQ 150MG BASE</u>	<u>A213796 001</u>	Apr 19, 2022
<u>AB</u>	AIPING PHARM INC	<u>EQ 10MG BASE</u>	<u>A213474 001</u>	Jul 28, 2020
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>A213474 002</u>	Jul 28, 2020
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A213474 003</u>	Jul 28, 2020
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A213474 004</u>	Jul 28, 2020
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A213474 005</u>	Jul 28, 2020
<u>AB</u>	AJANTA PHARMA LTD	<u>EQ 10MG BASE</u>	<u>A212624 001</u>	Sep 13, 2019
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>A212624 002</u>	Sep 13, 2019
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A212624 003</u>	Sep 13, 2019
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A212624 004</u>	Sep 13, 2019
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A212624 005</u>	Sep 13, 2019
<u>AB</u>	ALEMBIC	<u>EQ 10MG BASE</u>	<u>A215076 001</u>	Apr 21, 2021
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>A215076 002</u>	Apr 21, 2021
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A215076 003</u>	Apr 21, 2021
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A215076 004</u>	Apr 21, 2021
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A215076 005</u>	Apr 21, 2021
<u>AB</u>	AMNEAL PHARMS CO	<u>EQ 10MG BASE</u>	<u>A207482 001</u>	Jun 28, 2017
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>A207482 002</u>	Jun 28, 2017
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A207482 003</u>	Jun 28, 2017
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A207482 004</u>	Jun 28, 2017
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A207482 005</u>	Jun 28, 2017
<u>AB</u>	APPCO	<u>EQ 10MG BASE</u>	<u>A214908 001</u>	Apr 01, 2021
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>A214908 002</u>	Apr 01, 2021
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A214908 003</u>	Apr 01, 2021
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A214908 004</u>	Apr 01, 2021
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A214908 005</u>	Apr 01, 2021
<u>AB</u>	AUROBINDO PHARMA LTD	<u>EQ 10MG BASE</u>	<u>A211603 001</u>	Mar 27, 2019
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>A211603 002</u>	Mar 27, 2019
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A211603 003</u>	Mar 27, 2019
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A211603 004</u>	Mar 27, 2019
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A211603 005</u>	Mar 27, 2019
<u>AB</u>	CHARTWELL RX	<u>EQ 10MG BASE</u>	<u>A210268 001</u>	Sep 04, 2020
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>A210268 002</u>	Sep 04, 2020
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A210268 003</u>	Sep 04, 2020
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A210268 004</u>	Sep 04, 2020
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A210268 005</u>	Sep 04, 2020
<u>AB</u>	! ENDO OPERATIONS	<u>EQ 150MG BASE</u>	<u>A071422 006</u>	Nov 09, 1987
<u>AB</u>	EPIC PHARMA LLC	<u>EQ 10MG BASE</u>	<u>A210675 001</u>	Oct 16, 2020
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>A210675 002</u>	Oct 16, 2020
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A210675 004</u>	Mar 03, 2022
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A210675 005</u>	Mar 03, 2022
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A210675 003</u>	Oct 16, 2020
<u>AB</u>	JUBILANT CADISTA	<u>EQ 10MG BASE</u>	<u>A215483 001</u>	Mar 14, 2022
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>A215483 002</u>	Mar 14, 2022
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A215483 003</u>	Mar 14, 2022
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A215483 004</u>	Mar 14, 2022
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A215483 005</u>	Mar 14, 2022
<u>AB</u>	LEADING	<u>EQ 10MG BASE</u>	<u>A211619 001</u>	Mar 09, 2021
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>A211619 002</u>	Mar 09, 2021
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A211619 003</u>	Mar 09, 2021
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A211619 004</u>	Mar 09, 2021
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A211619 005</u>	Mar 09, 2021
<u>AB</u>	MANKIND PHARMA	<u>EQ 10MG BASE</u>	<u>A215710 001</u>	Feb 09, 2022

PRESCRIPTION DRUG PRODUCT LIST

DOXEPIN HYDROCHLORIDE

CAPSULE; ORAL

DOXEPIN HYDROCHLORIDE

<u>AB</u>		<u>EQ 25MG BASE</u>	<u>A215710 002</u>	Feb 09, 2022
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A215710 003</u>	Feb 09, 2022
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A215710 004</u>	Feb 09, 2022
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A215710 005</u>	Feb 09, 2022
<u>AB</u>	MICRO LABS	<u>EQ 10MG BASE</u>	<u>A217688 001</u>	Jun 27, 2023
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>A217688 002</u>	Jun 27, 2023
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A217688 003</u>	Jun 27, 2023
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A217688 004</u>	Jun 27, 2023
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A217688 005</u>	Jun 27, 2023
<u>AB</u>	MSN	<u>EQ 10MG BASE</u>	<u>A215113 001</u>	Jun 24, 2022
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>A215113 002</u>	Jun 24, 2022
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A215113 003</u>	Jun 24, 2022
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A215113 004</u>	Jun 24, 2022
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A215113 005</u>	Jun 24, 2022
<u>AB</u>	MYLAN PHARMS INC	<u>EQ 10MG BASE</u>	<u>A070791 002</u>	May 13, 1986
<u>AB</u>	!	<u>EQ 25MG BASE</u>	<u>A070791 003</u>	May 13, 1986
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A070791 001</u>	May 13, 1986
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A070791 004</u>	May 13, 1986
<u>AB</u>	!	<u>EQ 100MG BASE</u>	<u>A070791 005</u>	May 13, 1986
<u>AB</u>	TARO	<u>EQ 10MG BASE</u>	<u>A213063 001</u>	Jul 01, 2020
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>A213063 002</u>	Jul 01, 2020
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A213063 003</u>	Jul 01, 2020
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A213063 004</u>	Jul 01, 2020
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A213063 005</u>	Jul 01, 2020
<u>AB</u>	UNIQUE PHARM	<u>EQ 10MG BASE</u>	<u>A217975 001</u>	Aug 21, 2023
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>A217975 002</u>	Aug 21, 2023
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A217975 003</u>	Aug 21, 2023
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A217975 004</u>	Aug 21, 2023
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A217975 005</u>	Aug 21, 2023
<u>AB</u>	VELZEN PHARMA PVT	<u>EQ 150MG BASE</u>	<u>A211618 001</u>	Mar 01, 2021
<u>AB</u>	ZYDUS LIFESCIENCES	<u>EQ 10MG BASE</u>	<u>A210700 001</u>	Mar 23, 2023
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>A210700 002</u>	Mar 23, 2023
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A210700 003</u>	Mar 23, 2023
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A210700 004</u>	Mar 23, 2023
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A210700 005</u>	Mar 23, 2023
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A210140 001</u>	Mar 21, 2023

CONCENTRATE; ORAL

DOXEPIN HYDROCHLORIDE

! LANNETT CO INC

EQ 10MG BASE/ML

A074721 001 Dec 29, 1998

CREAM; TOPICAL

DOXEPIN HYDROCHLORIDE

<u>AB</u>	AMNEAL	<u>5%</u>	<u>A212357 001</u>	Aug 16, 2023
<u>AB</u>	TEVA PHARMS	<u>5%</u>	<u>A215408 001</u>	Feb 17, 2023
<u>AB</u>	+	<u>5%</u>	<u>N020126 001</u>	Apr 01, 1994

TABLET; ORAL

DOXEPIN HYDROCHLORIDE

<u>AB</u>	ACTAVIS ELIZABETH	<u>EQ 3MG BASE</u>	<u>A201951 001</u>	Jul 26, 2013
<u>AB</u>		<u>EQ 6MG BASE</u>	<u>A201951 002</u>	Jul 26, 2013
<u>AB</u>	MSN	<u>EQ 3MG BASE</u>	<u>A214823 001</u>	Apr 03, 2023
<u>AB</u>		<u>EQ 6MG BASE</u>	<u>A214823 002</u>	Apr 03, 2023
<u>AB</u>	RK PHARMA	<u>EQ 3MG BASE</u>	<u>A202337 001</u>	Jan 20, 2016
<u>AB</u>		<u>EQ 6MG BASE</u>	<u>A202337 002</u>	Jan 20, 2016
<u>AB</u>	STRIDES PHARMA	<u>EQ 3MG BASE</u>	<u>A202510 001</u>	Jul 24, 2020
<u>AB</u>		<u>EQ 6MG BASE</u>	<u>A202510 002</u>	Jul 24, 2020
<u>AB</u>	ZYDUS LIFESCIENCES	<u>EQ 3MG BASE</u>	<u>A202761 001</u>	Aug 16, 2023
<u>AB</u>		<u>EQ 6MG BASE</u>	<u>A202761 002</u>	Aug 16, 2023
<u>AB</u>	+	<u>EQ 3MG BASE</u>	<u>N022036 001</u>	Mar 17, 2010
<u>AB</u>	+	<u>EQ 6MG BASE</u>	<u>N022036 002</u>	Mar 17, 2010

DOXERCALCIFEROL

CAPSULE; ORAL

DOXERCALCIFEROL

<u>AB</u>	AVET	<u>0.5MCG</u>	<u>A205360 001</u>	Sep 15, 2020
<u>AB</u>		<u>1MCG</u>	<u>A205360 002</u>	Sep 15, 2020
<u>AB</u>		<u>2.5MCG</u>	<u>A205360 003</u>	Sep 15, 2020
<u>AB</u>	RISING	<u>0.5MCG</u>	<u>A201518 001</u>	Sep 09, 2016
<u>AB</u>		<u>1MCG</u>	<u>A201518 002</u>	Sep 09, 2016
<u>AB</u>		<u>2.5MCG</u>	<u>A201518 003</u>	Sep 09, 2016

PRESCRIPTION DRUG PRODUCT LIST

DOXERCALCIFEROL

CAPSULE; ORAL

HECTOROL

<u>AB</u>	+	SANOFI	<u>0.5MCG</u>	<u>N020862</u>	<u>002</u>	Apr 23, 2004
<u>AB</u>	+		<u>1MCG</u>	<u>N020862</u>	<u>003</u>	Jul 13, 2009
<u>AB</u>	+	!	<u>2.5MCG</u>	<u>N020862</u>	<u>001</u>	Jun 09, 1999

INJECTABLE; INJECTION

DOXERCALCIFEROL

<u>AP</u>		ALEMBIC	<u>4MCG/2ML (2MCG/ML)</u>	<u>A215810</u>	<u>001</u>	Jun 15, 2023
<u>AP</u>		EUGIA PHARMA	<u>2MCG/ML (2MCG/ML)</u>	<u>A213717</u>	<u>001</u>	Jan 24, 2022
<u>AP</u>			<u>4MCG/2ML (2MCG/ML)</u>	<u>A213717</u>	<u>002</u>	Jan 24, 2022
<u>AP</u>		GLAND PHARMA LTD	<u>4MCG/2ML (2MCG/ML)</u>	<u>A210452</u>	<u>001</u>	Sep 26, 2019
<u>AP</u>		HIKMA	<u>4MCG/2ML (2MCG/ML)</u>	<u>A091101</u>	<u>001</u>	Aug 30, 2013
<u>AP</u>	+	HOSPIRA	<u>4MCG/2ML (2MCG/ML)</u>	<u>N208614</u>	<u>001</u>	Jul 24, 2018
<u>AP</u>		LUPIN LTD	<u>4MCG/2ML (2MCG/ML)</u>	<u>A210801</u>	<u>001</u>	Nov 01, 2018
<u>AP</u>		MEITHEAL	<u>4MCG/2ML (2MCG/ML)</u>	<u>A211670</u>	<u>001</u>	Feb 07, 2020
<u>AP</u>		SANDOZ	<u>4MCG/2ML (2MCG/ML)</u>	<u>A091333</u>	<u>001</u>	May 05, 2014
<u>AP</u>			<u>4MCG/2ML (2MCG/ML)</u>	<u>A200926</u>	<u>001</u>	Feb 04, 2014

HECTOROL

<u>AP</u>	+	SANOFI	<u>2MCG/ML (2MCG/ML)</u>	<u>N021027</u>	<u>002</u>	Apr 06, 2000
<u>AP</u>	+	!	<u>4MCG/2ML (2MCG/ML)</u>	<u>N021027</u>	<u>001</u>	Apr 06, 2000

DOXORUBICIN HYDROCHLORIDE

INJECTABLE; INJECTION

DOXORUBICIN HYDROCHLORIDE

<u>AP</u>		ACTAVIS INC	<u>2MG/ML</u>	<u>A203622</u>	<u>001</u>	Jun 27, 2014
<u>AP</u>			<u>200MG/100ML</u>	<u>A203622</u>	<u>002</u>	Jun 27, 2014
<u>AP</u>		AMNEAL	<u>20MG/VIAL</u>	<u>A208888</u>	<u>001</u>	Feb 17, 2017
<u>AP</u>			<u>50MG/VIAL</u>	<u>A208888</u>	<u>002</u>	Feb 17, 2017
<u>AP</u>		FRESENIUS KABI USA	<u>2MG/ML</u>	<u>A063277</u>	<u>001</u>	Oct 26, 1995
<u>AP</u>		GLAND PHARMA LTD	<u>2MG/ML</u>	<u>A209825</u>	<u>001</u>	Aug 11, 2017
<u>AP</u>		HIKMA	<u>2MG/ML</u>	<u>A062975</u>	<u>001</u>	Mar 17, 1989
<u>AP</u>	!		<u>20MG/VIAL</u>	<u>A062921</u>	<u>002</u>	Mar 17, 1989
<u>AP</u>	!		<u>50MG/VIAL</u>	<u>A062921</u>	<u>003</u>	Mar 17, 1989
<u>AP</u>			<u>200MG/100ML</u>	<u>A064097</u>	<u>001</u>	Sep 13, 1994
<u>AP</u>		MYLAN LABS LTD	<u>50MG/VIAL</u>	<u>A200170</u>	<u>002</u>	Oct 28, 2011
<u>AP</u>	+	PFIZER	<u>2MG/ML</u>	<u>N050629</u>	<u>001</u>	Dec 23, 1987
<u>AP</u>	+	!	<u>200MG/100ML</u>	<u>N050629</u>	<u>002</u>	May 03, 1988
<u>AP</u>		SAGENT PHARMS	<u>2MG/ML</u>	<u>A091495</u>	<u>001</u>	Mar 18, 2013
<u>AP</u>		SUN PHARM INDS	<u>2MG/ML</u>	<u>A091418</u>	<u>001</u>	Feb 15, 2012
<u>AP</u>	!	HIKMA	<u>10MG/VIAL</u>	<u>A062921</u>	<u>001</u>	Mar 17, 1989

INJECTABLE, LIPOSOMAL; INJECTION

DOXIL (LIPOSOMAL)

<u>AB</u>	+	BAXTER HLTHCARE CORP	<u>20MG/10ML (2MG/ML)</u>	<u>N050718</u>	<u>001</u>	Nov 17, 1995
<u>AB</u>	+		<u>50MG/25ML (2MG/ML)</u>	<u>N050718</u>	<u>002</u>	Jun 13, 2000

DOXORUBICIN HYDROCHLORIDE (LIPOSOMAL)

<u>AB</u>		AYANA PHARMA LTD	<u>20MG/10ML (2MG/ML)</u>	<u>A207228</u>	<u>001</u>	Oct 12, 2021
<u>AB</u>			<u>50MG/25ML (2MG/ML)</u>	<u>A207228</u>	<u>002</u>	Oct 12, 2021
<u>AB</u>		BAXTER HLTHCARE CORP	<u>20MG/10ML (2MG/ML)</u>	<u>A212219</u>	<u>001</u>	Oct 19, 2022
<u>AB</u>			<u>50MG/25ML (2MG/ML)</u>	<u>A212219</u>	<u>002</u>	Oct 19, 2022
<u>AB</u>		DR REDDYS	<u>20MG/10ML (2MG/ML)</u>	<u>A208657</u>	<u>001</u>	May 15, 2017
<u>AB</u>			<u>50MG/25ML (2MG/ML)</u>	<u>A208657</u>	<u>002</u>	May 15, 2017
<u>AB</u>		LUPIN	<u>20MG/10ML (2MG/ML)</u>	<u>A215178</u>	<u>001</u>	Jul 16, 2024
<u>AB</u>			<u>50MG/25ML (2MG/ML)</u>	<u>A215178</u>	<u>002</u>	Jul 16, 2024
<u>AB</u>	!	SUN PHARM	<u>20MG/10ML (2MG/ML)</u>	<u>A203263</u>	<u>001</u>	Feb 04, 2013
<u>AB</u>	!		<u>50MG/25ML (2MG/ML)</u>	<u>A203263</u>	<u>002</u>	Feb 04, 2013
<u>AB</u>		ZYDUS LIFESCIENCES	<u>20MG/10ML (2MG/ML)</u>	<u>A212299</u>	<u>001</u>	Sep 10, 2020
<u>AB</u>			<u>50MG/25ML (2MG/ML)</u>	<u>A212299</u>	<u>002</u>	Sep 10, 2020

DOXYCYCLINE

CAPSULE; ORAL

DOXYCYCLINE

<u>AB</u>		ALEMBIC	<u>40MG</u>	<u>A217674</u>	<u>001</u>	Jun 27, 2024
<u>AB</u>			<u>EQ 75MG BASE</u>	<u>A209165</u>	<u>001</u>	Jul 28, 2017
<u>AB</u>			<u>EQ 100MG BASE</u>	<u>A209165</u>	<u>002</u>	Jul 28, 2017
<u>AB</u>		APOTEX	<u>40MG</u>	<u>A217170</u>	<u>001</u>	Jul 11, 2024
<u>AB</u>		DR REDDYS	<u>40MG</u>	<u>A218034</u>	<u>001</u>	Apr 08, 2024
<u>AB</u>		DR REDDYS LABS SA	<u>EQ 50MG BASE</u>	<u>A209396</u>	<u>001</u>	Sep 29, 2017
<u>AB</u>			<u>EQ 75MG BASE</u>	<u>A209396</u>	<u>002</u>	Sep 29, 2017
<u>AB</u>			<u>EQ 100MG BASE</u>	<u>A209396</u>	<u>003</u>	Sep 29, 2017
<u>AB</u>		LUPIN	<u>40MG</u>	<u>A216631</u>	<u>001</u>	Apr 08, 2024
<u>AB</u>		LUPIN LTD	<u>EQ 50MG BASE</u>	<u>A204234</u>	<u>001</u>	Mar 05, 2014

PRESCRIPTION DRUG PRODUCT LIST

DOXYCYCLINE

CAPSULE; ORAL

DOXYCYCLINE

<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A204234 002</u>	Mar 05, 2014
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A204234 003</u>	Mar 05, 2014
<u>AB</u>	STRIDES PHARMA	<u>EQ 50MG BASE</u>	<u>A065055 001</u>	Dec 01, 2000
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A065055 002</u>	Dec 01, 2000
<u>AB</u>	!	<u>EQ 150MG BASE</u>	<u>A065055 003</u>	Jul 15, 2005
<u>AB</u>	SUN PHARM INDS LTD	<u>EQ 50MG BASE</u>	<u>A065053 001</u>	Nov 22, 2000
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A065053 003</u>	Sep 10, 2003
<u>AB</u>	!	<u>EQ 100MG BASE</u>	<u>A065053 002</u>	Nov 22, 2000
<u>AB</u>	ZYDUS PHARMS	<u>EQ 50MG BASE</u>	<u>A205115 001</u>	Feb 18, 2016
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A205115 002</u>	Feb 18, 2016
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A205115 003</u>	Feb 18, 2016

MONODOX

<u>AB</u>	+	CHARTWELL RX	<u>EQ 50MG BASE</u>	<u>N050641 002</u>	Feb 10, 1992
<u>AB</u>	+		<u>EQ 75MG BASE</u>	<u>N050641 003</u>	Oct 18, 2006
<u>AB</u>	+		<u>EQ 100MG BASE</u>	<u>N050641 001</u>	Dec 29, 1989
<u>AB</u>	+		<u>EQ 150MG BASE</u>	<u>N050641 004</u>	Feb 17, 2022

ORACEA

<u>AB</u>	+	!	GALDERMA LABS LP	<u>40MG</u>	<u>N050805 001</u>	May 26, 2006
-----------	---	---	------------------	-------------	--------------------	--------------

FOR SUSPENSION; ORAL

DOXYCYCLINE

<u>AB</u>		CHARTWELL	<u>EQ 25MG BASE/5ML</u>	<u>A065454 001</u>	Jul 16, 2008
<u>AB</u>	!	LUPIN LTD	<u>EQ 25MG BASE/5ML</u>	<u>A201678 001</u>	Mar 18, 2013

TABLET; ORAL

DOXYCYCLINE

<u>AB</u>		HERITAGE	<u>EQ 50MG BASE</u>	<u>A091605 001</u>	Dec 20, 2011
<u>AB</u>			<u>EQ 75MG BASE</u>	<u>A091605 002</u>	Dec 20, 2011
<u>AB</u>			<u>EQ 100MG BASE</u>	<u>A091605 003</u>	Dec 20, 2011
<u>AB</u>	!		<u>EQ 150MG BASE</u>	<u>A091605 004</u>	Dec 20, 2011
<u>AB</u>		LANNETT CO INC	<u>EQ 50MG BASE</u>	<u>A065285 001</u>	Dec 08, 2005
<u>AB</u>			<u>EQ 75MG BASE</u>	<u>A065285 003</u>	Jul 30, 2008
<u>AB</u>			<u>EQ 100MG BASE</u>	<u>A065285 002</u>	Dec 08, 2005
<u>AB</u>			<u>EQ 150MG BASE</u>	<u>A065285 004</u>	Jul 30, 2008
<u>AB</u>		STRIDES PHARMA	<u>EQ 50MG BASE</u>	<u>A065070 001</u>	Dec 15, 2000
<u>AB</u>			<u>EQ 75MG BASE</u>	<u>A065070 003</u>	Dec 30, 2002
<u>AB</u>			<u>EQ 100MG BASE</u>	<u>A065070 002</u>	Dec 15, 2000
<u>AB</u>			<u>EQ 150MG BASE</u>	<u>A065070 004</u>	Jul 14, 2005
<u>AB</u>		SUN PHARM INDS LTD	<u>EQ 50MG BASE</u>	<u>A065356 001</u>	May 31, 2006
<u>AB</u>			<u>EQ 75MG BASE</u>	<u>A065356 002</u>	May 31, 2006
<u>AB</u>			<u>EQ 100MG BASE</u>	<u>A065356 003</u>	May 31, 2006
<u>AB</u>			<u>EQ 150MG BASE</u>	<u>A065356 004</u>	Jul 29, 2010
<u>AB</u>		ZYDUS PHARMS	<u>EQ 50MG BASE</u>	<u>A209582 001</u>	Sep 28, 2017
<u>AB</u>			<u>EQ 75MG BASE</u>	<u>A209582 002</u>	Sep 28, 2017
<u>AB</u>			<u>EQ 100MG BASE</u>	<u>A209582 003</u>	Sep 28, 2017
<u>AB</u>			<u>EQ 150MG BASE</u>	<u>A209582 004</u>	Sep 28, 2017

DOXYCYCLINE HYCLATE

CAPSULE; ORAL

DOXYCYCLINE HYCLATE

<u>AB</u>		ACTAVIS LABS FL INC	<u>EQ 50MG BASE</u>	<u>A062031 002</u>	Oct 13, 1982
<u>AB</u>			<u>EQ 100MG BASE</u>	<u>A062031 001</u>	
<u>AB</u>		ALEMBIC	<u>EQ 50MG BASE</u>	<u>A210527 001</u>	Jun 13, 2018
<u>AB</u>			<u>EQ 100MG BASE</u>	<u>A210527 002</u>	Jun 13, 2018
<u>AB</u>		AMNEAL PHARMS	<u>EQ 100MG BASE</u>	<u>A207289 001</u>	Jun 27, 2016
<u>AB</u>		CHANGZHOU PHARM	<u>EQ 100MG BASE</u>	<u>A209402 001</u>	Oct 07, 2019
<u>AB</u>		CHARTWELL	<u>EQ 50MG BASE</u>	<u>A062500 001</u>	Sep 11, 1984
<u>AB</u>			<u>EQ 100MG BASE</u>	<u>A062500 002</u>	Sep 11, 1984
<u>AB</u>		HIKMA INTL PHARMS	<u>EQ 50MG BASE</u>	<u>A062396 002</u>	Nov 07, 1984
<u>AB</u>			<u>EQ 100MG BASE</u>	<u>A062396 001</u>	May 07, 1984
<u>AB</u>		SUN PHARM INDUSTRIES	<u>EQ 50MG BASE</u>	<u>A062676 002</u>	Jul 10, 1986
<u>AB</u>			<u>EQ 100MG BASE</u>	<u>A062676 001</u>	Jul 10, 1986
<u>AB</u>		ZYDUS LIFESCIENCES	<u>EQ 50MG BASE</u>	<u>A207774 001</u>	May 31, 2018
<u>AB</u>			<u>EQ 100MG BASE</u>	<u>A207774 002</u>	May 31, 2018

VIBRAMYCIN

<u>AB</u>	+	!	PFIZER	<u>EQ 100MG BASE</u>	<u>N050007 002</u>
-----------	---	---	--------	----------------------	--------------------

INJECTABLE; INJECTION

DOXY 100

<u>AP</u>	!	FRESENIUS KABI USA	<u>EQ 100MG BASE/VIAL</u>	<u>A062475 001</u>	Dec 09, 1983
-----------	---	--------------------	---------------------------	--------------------	--------------

DOXY 200

<u>AP</u>	!	FRESENIUS KABI USA	<u>EQ 200MG BASE/VIAL</u>	<u>A062475 002</u>	Dec 09, 1983
-----------	---	--------------------	---------------------------	--------------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

DOXYCYCLINE HYCLATE

INJECTABLE; INJECTION

DOXYCYCLINE

<u>AP</u>	MYLAN LABS LTD	<u>EQ 100MG BASE/VIAL</u>	<u>A091406 001</u>	Aug 21, 2012
<u>AP</u>	! ZYDUS PHARMS	<u>EQ 100MG BASE/VIAL</u>	<u>A207757 001</u>	Sep 28, 2017
<u>AP</u>		<u>EQ 200MG BASE/VIAL</u>	<u>A207757 002</u>	Sep 28, 2017

DOXYCYCLINE HYCLATE

<u>AP</u>	AMNEAL	<u>EQ 100MG BASE/VIAL</u>	<u>A217487 001</u>	Dec 04, 2024
<u>AP</u>	ENDO OPERATIONS	<u>EQ 100MG BASE/VIAL</u>	<u>A216690 001</u>	Dec 07, 2022
<u>AP</u>	GLAND PHARMA LTD	<u>EQ 100MG BASE/VIAL</u>	<u>A215583 001</u>	Apr 12, 2023
<u>AP</u>	HERITAGE	<u>EQ 100MG BASE/VIAL</u>	<u>A217854 001</u>	Jul 23, 2024
<u>AP</u>	KINDOS	<u>EQ 100MG BASE/VIAL</u>	<u>A218053 001</u>	Oct 28, 2024
<u>AP</u>	LUPIN LTD	<u>EQ 100MG BASE/VIAL</u>	<u>A217794 001</u>	Mar 14, 2024
<u>AP</u>	SLATE RUN PHARMA	<u>EQ 100MG BASE/VIAL</u>	<u>A217685 001</u>	Jun 21, 2024
<u>AP</u>		<u>EQ 200MG BASE/VIAL</u>	<u>A217685 002</u>	Jun 21, 2024
<u>AP</u>	WEST-WARD PHARMS INT	<u>EQ 100MG BASE/VIAL</u>	<u>A062992 001</u>	Feb 16, 1989
<u>AP</u>		<u>EQ 200MG BASE/VIAL</u>	<u>A062992 002</u>	Feb 16, 1989

TABLET; ORAL

ACTICLATE

<u>AB</u>	+ CHARTWELL RX	<u>EQ 75MG BASE</u>	<u>N205931 001</u>	Jul 25, 2014
<u>AB</u>	+!	<u>EQ 150MG BASE</u>	<u>N205931 002</u>	Jul 25, 2014

DOXYCYCLINE HYCLATE

<u>AB</u>	ACELLA	<u>EQ 100MG BASE</u>	<u>A210664 001</u>	Mar 16, 2020
<u>AB</u>	ACTAVIS LABS FL INC	<u>EQ 100MG BASE</u>	<u>A062421 001</u>	Feb 02, 1983
<u>AB</u>	AJANTA PHARMA LTD	<u>EQ 75MG BASE</u>	<u>A211584 001</u>	Jun 01, 2020
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A211584 002</u>	Jun 01, 2020
<u>AB</u>	ALEMBIC	<u>EQ 20MG BASE</u>	<u>A210537 001</u>	Mar 03, 2020
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A210536 002</u>	Sep 14, 2021
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A211744 001</u>	Jun 30, 2020
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A211744 002</u>	Jun 30, 2020
<u>AB</u>	APOTEX	<u>EQ 75MG BASE</u>	<u>A209243 001</u>	Apr 15, 2019
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A209243 002</u>	Apr 15, 2019
<u>AB</u>	CHANGZHOU PHARM	<u>EQ 100MG BASE</u>	<u>A211343 001</u>	Oct 09, 2019
<u>AB</u>	CHARTWELL	<u>EQ 50MG BASE</u>	<u>A062505 002</u>	Jul 27, 2021
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A062505 001</u>	Sep 11, 1984
<u>AB</u>	! CHARTWELL MOLECULAR	<u>EQ 20MG BASE</u>	<u>A065277 001</u>	Nov 10, 2005
<u>AB</u>	DR REDDYS LABS SA	<u>EQ 75MG BASE</u>	<u>A208765 001</u>	Jun 14, 2017
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A208765 002</u>	Jun 14, 2017
<u>AB</u>	EPIC PHARMA LLC	<u>EQ 20MG BASE</u>	<u>A065182 001</u>	May 13, 2005
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A062269 003</u>	Oct 05, 1983
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A214207 001</u>	Dec 16, 2020
<u>AB</u>	!	<u>EQ 100MG BASE</u>	<u>A062269 002</u>	Nov 08, 1982
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A214207 002</u>	Dec 16, 2020
<u>AB</u>	HIKMA INTL PHARMS	<u>EQ 100MG BASE</u>	<u>A065095 001</u>	Jul 02, 2003
<u>AB</u>	LARKEN LABS	<u>EQ 20MG BASE</u>	<u>A065287 001</u>	Feb 28, 2006
<u>AB</u>	LUPIN LTD	<u>EQ 75MG BASE</u>	<u>A208818 001</u>	Sep 27, 2017
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A208818 002</u>	Sep 27, 2017
<u>AB</u>	MYLAN	<u>EQ 100MG BASE</u>	<u>A062432 001</u>	Feb 15, 1983
<u>AB</u>	NOVEL LABS INC	<u>EQ 100MG BASE</u>	<u>A207558 001</u>	Sep 06, 2017
<u>AB</u>	SUN PHARM INDUSTRIES	<u>EQ 20MG BASE</u>	<u>A065134 001</u>	May 13, 2005
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A062677 001</u>	Jul 10, 1986
<u>AB</u>	ZYDUS LIFESCIENCES	<u>EQ 100MG BASE</u>	<u>A207773 001</u>	Oct 30, 2017

TABLET, DELAYED RELEASE; ORAL

DORYX

<u>AB</u>	+ MAYNE PHARMA	<u>EQ 200MG BASE</u>	<u>N050795 005</u>	Apr 11, 2013
-----------	----------------	----------------------	--------------------	--------------

DOXYCYCLINE HYCLATE

<u>AB</u>	ACTAVIS ELIZABETH	<u>EQ 50MG BASE</u>	<u>A090134 003</u>	May 22, 2018
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A090134 001</u>	Dec 14, 2011
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A090134 002</u>	Dec 14, 2011
<u>AB</u>		<u>EQ 200MG BASE</u>	<u>A090134 004</u>	May 22, 2018
<u>AB</u>	ALEMBIC	<u>EQ 75MG BASE</u>	<u>A213075 001</u>	Jan 03, 2022
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A213075 002</u>	Jan 03, 2022
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A213075 003</u>	Jan 03, 2022
<u>AB</u>		<u>EQ 200MG BASE</u>	<u>A213075 004</u>	Jan 03, 2022
<u>AB</u>	HERITAGE	<u>EQ 75MG BASE</u>	<u>A200856 001</u>	Apr 30, 2013
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A200856 002</u>	Apr 30, 2013
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A200856 003</u>	Apr 30, 2013
<u>AB</u>	!	<u>EQ 200MG BASE</u>	<u>A200856 004</u>	Nov 13, 2018
<u>AB</u>	PRINSTON INC	<u>EQ 50MG BASE</u>	<u>A207494 003</u>	Feb 19, 2019
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A207494 001</u>	Nov 15, 2016
<u>AB</u>		<u>EQ 200MG BASE</u>	<u>A207494 002</u>	Nov 15, 2016

PRESCRIPTION DRUG PRODUCT LIST

DOXYCYCLINE HYCLATE

TABLET, DELAYED RELEASE;ORAL

DORYX

+ MAYNE PHARMA EQ 80MG BASE

N050795 004 Apr 11, 2013

DORYX MPC

+ MAYNE PHARMA EQ 60MG BASE

N050795 007 May 20, 2016

DOXYLAMINE SUCCINATE; PYRIDOXINE HYDROCHLORIDE

TABLET, DELAYED RELEASE;ORAL

DICLEGIS**AB** +! DUCHESNAY **10MG;10MG****N021876 001** Apr 08, 2013DOXYLAMINE SUCCINATE AND PYRIDOXINE HYDROCHLORIDE**AB** ACTAVIS LABS FL INC **10MG;10MG****A205811 001** Aug 19, 2016**AB** BIONPHARMA **10MG;10MG****A217000 001** Aug 04, 2023**AB** ENDO OPERATIONS **10MG;10MG****A208518 001** Dec 06, 2017**AB** MYLAN PHARMS INC **10MG;10MG****A207825 001** Jul 06, 2020

TABLET, EXTENDED RELEASE;ORAL

BONJESTA

+! DUCHESNAY 20MG;20MG

N209661 001 Nov 07, 2016

DRONABINOL

CAPSULE;ORAL

DRONABINOL**AB** ASCENT PHARMS INC **2.5MG****A207421 001** Feb 07, 2020**AB** **5MG****A207421 002** Feb 07, 2020**AB** **10MG****A207421 003** Feb 07, 2020MARINOL**AB** + ALKEM LABS LTD **2.5MG****N018651 001** May 31, 1985**AB** +! **5MG****N018651 002** May 31, 1985**AB** + **10MG****N018651 003** May 31, 1985DRONEDARONE HYDROCHLORIDE

TABLET;ORAL

DRONEDARONE HYDROCHLORIDE**AB** LUPIN **EQ 400MG BASE****A205904 001** Jan 31, 2024MULTAQ**AB** +! SANOFI AVENTIS US **EQ 400MG BASE****N022425 001** Jul 01, 2009DROPERIDOL

INJECTABLE; INJECTION

DROPERIDOL**AP** ! AM REGENT **2.5MG/ML****A072123 001** Oct 24, 1988**AP** HIKMA **2.5MG/ML****A208197 001** Dec 14, 2017DROSPIRENONE

TABLET;ORAL

SLYND

+! EXELTIS USA INC 4MG

N211367 001 May 23, 2019

DROSPIRENONE; ESTETROL

TABLET;ORAL

NEXTSTELLIS

+! MAYNE PHARMA 3MG;14.2MG

N214154 001 Apr 15, 2021

DROSPIRENONE; ESTRADIOL

TABLET;ORAL

ANGELIQ

+ BAYER HLTHCARE 0.25MG;0.5MG

N021355 001 Feb 29, 2012

+! 0.5MG;1MG

N021355 002 Sep 28, 2005

DROSPIRENONE; ETHINYL ESTRADIOL

TABLET;ORAL

DROSPIRENONE AND ETHINYL ESTRADIOL**AB** GLENMARK PHARMS LTD **3MG;0.02MG****A204296 001** Aug 17, 2015**AB** HETERO LABS **3MG;0.02MG****A211944 001** Mar 22, 2019**AB** HLTHCARE **3MG;0.02MG****A203291 001** Jul 18, 2017**AB** WATSON LABS **3MG;0.02MG****A078833 001** Nov 28, 2011**AB** XIROMED **3MG;0.02MG****A202594 001** Oct 22, 2015LO-ZUMANDIMINE**AB** AUROBINDO PHARMA **3MG;0.02MG****A209632 001** Feb 27, 2018

LTD

LORYNA**AB** XIROMED **3MG;0.02MG****A079221 001** Mar 28, 2011MELAMISA**AB** NOVAST LABS **3MG;0.02MG****A202016 001** Jan 26, 2016

PRESCRIPTION DRUG PRODUCT LIST

DROSPIRENONE; ETHINYL ESTRADIOL

TABLET; ORAL

NIKKI

<u>AB</u>	LUPIN LTD	<u>3MG;0.02MG</u>	<u>A201661 001</u>	May 27, 2014
-----------	-----------	-------------------	--------------------	--------------

YAZ

<u>AB</u>	+! BAYER HLTHCARE	<u>3MG;0.02MG</u>	<u>N021676 001</u>	Mar 16, 2006
-----------	-------------------	-------------------	--------------------	--------------

TABLET; ORAL-28

DROSPIRENONE AND ETHINYL ESTRADIOL

<u>AB</u>	DR REDDYS LABS SA	<u>3MG;0.03MG</u>	<u>A090081 001</u>	Sep 07, 2010
-----------	-------------------	-------------------	--------------------	--------------

<u>AB</u>	GLENMARK PHARMS LTD	<u>3MG;0.03MG</u>	<u>A204848 001</u>	Mar 25, 2016
-----------	---------------------	-------------------	--------------------	--------------

<u>AB</u>	HETERO LABS	<u>3MG;0.03MG</u>	<u>A213034 001</u>	Jan 24, 2020
-----------	-------------	-------------------	--------------------	--------------

<u>AB</u>	LUPIN LTD	<u>3MG;0.03MG</u>	<u>A201663 001</u>	Dec 18, 2012
-----------	-----------	-------------------	--------------------	--------------

<u>AB</u>	NAARI PTE LTD	<u>3MG;0.03MG</u>	<u>A207245 001</u>	Nov 22, 2016
-----------	---------------	-------------------	--------------------	--------------

<u>AB</u>	XIROMED	<u>3MG;0.03MG</u>	<u>A202131 001</u>	May 04, 2015
-----------	---------	-------------------	--------------------	--------------

SYEDA

<u>AB</u>	XIROMED	<u>3MG;0.03MG</u>	<u>A090114 001</u>	Mar 28, 2011
-----------	---------	-------------------	--------------------	--------------

YAELA

<u>AB</u>	NOVAST LABS	<u>3MG;0.03MG</u>	<u>A202015 001</u>	Nov 19, 2014
-----------	-------------	-------------------	--------------------	--------------

YASMIN

<u>AB</u>	+! BAYER HLTHCARE	<u>3MG;0.03MG</u>	<u>N021098 001</u>	May 11, 2001
-----------	-------------------	-------------------	--------------------	--------------

ZUMANDIMINE

<u>AB</u>	AUROBINDO PHARMA LTD	<u>3MG;0.03MG</u>	<u>A209407 001</u>	Mar 26, 2018
-----------	----------------------	-------------------	--------------------	--------------

DROSPIRENONE; ETHINYL ESTRADIOL; LEVOMEFOLATE CALCIUM

TABLET; ORAL

BEYAZ

<u>AB</u>	+! BAYER HLTHCARE	<u>3MG,N/A:0.02MG,N/A:0.451MG,0.451MG</u>	<u>N022532 001</u>	Sep 24, 2010
-----------	-------------------	---	--------------------	--------------

DROSPIRENONE, ETHINYL ESTRADIOL AND LEVOMEFOLATE CALCIUM

<u>AB</u>	LUPIN LTD	<u>3MG,N/A:0.02MG,N/A:0.451MG,0.451MG</u>	<u>A205947 001</u>	Jun 13, 2018
-----------	-----------	---	--------------------	--------------

<u>AB</u>	WATSON LABS INC	<u>3MG,N/A:0.02MG,N/A:0.451MG,0.451MG</u>	<u>A203593 001</u>	Oct 11, 2016
-----------	-----------------	---	--------------------	--------------

<u>AB</u>		<u>3MG,N/A:0.03MG,N/A:0.451MG,0.451MG</u>	<u>A203594 001</u>	Oct 11, 2016
-----------	--	---	--------------------	--------------

SAFYRAL

<u>AB</u>	+! BAYER HLTHCARE	<u>3MG,N/A:0.03MG,N/A:0.451MG,0.451MG</u>	<u>N022574 001</u>	Dec 16, 2010
-----------	-------------------	---	--------------------	--------------

TYDEMY

<u>AB</u>	LUPIN LTD	<u>3MG,N/A:0.03MG,N/A:0.451MG,0.451MG</u>	<u>A205948 001</u>	Dec 12, 2017
-----------	-----------	---	--------------------	--------------

DROXIDOPA

CAPSULE; ORAL

DROXIDOPA

<u>AB</u>	ADAPTIS	<u>100MG</u>	<u>A215265 001</u>	Nov 01, 2021
-----------	---------	--------------	--------------------	--------------

<u>AB</u>		<u>200MG</u>	<u>A215265 002</u>	Nov 01, 2021
-----------	--	--------------	--------------------	--------------

<u>AB</u>		<u>300MG</u>	<u>A215265 003</u>	Nov 01, 2021
-----------	--	--------------	--------------------	--------------

<u>AB</u>	AJANTA PHARMA LTD	<u>100MG</u>	<u>A214391 001</u>	Feb 18, 2021
-----------	-------------------	--------------	--------------------	--------------

<u>AB</u>		<u>200MG</u>	<u>A214391 002</u>	Feb 18, 2021
-----------	--	--------------	--------------------	--------------

<u>AB</u>		<u>300MG</u>	<u>A214391 003</u>	Feb 18, 2021
-----------	--	--------------	--------------------	--------------

<u>AB</u>	ALKEM LABS LTD	<u>100MG</u>	<u>A213911 001</u>	Feb 18, 2021
-----------	----------------	--------------	--------------------	--------------

<u>AB</u>		<u>200MG</u>	<u>A213911 002</u>	Feb 18, 2021
-----------	--	--------------	--------------------	--------------

<u>AB</u>		<u>300MG</u>	<u>A213911 003</u>	Feb 18, 2021
-----------	--	--------------	--------------------	--------------

<u>AB</u>	ANNORA	<u>100MG</u>	<u>A211726 002</u>	Aug 09, 2021
-----------	--------	--------------	--------------------	--------------

<u>AB</u>		<u>200MG</u>	<u>A211726 003</u>	Aug 09, 2021
-----------	--	--------------	--------------------	--------------

<u>AB</u>		<u>300MG</u>	<u>A211726 001</u>	Feb 18, 2021
-----------	--	--------------	--------------------	--------------

<u>AB</u>	AUROBINDO PHARMA LTD	<u>100MG</u>	<u>A214387 001</u>	Feb 18, 2021
-----------	----------------------	--------------	--------------------	--------------

<u>AB</u>		<u>200MG</u>	<u>A214387 002</u>	Feb 18, 2021
-----------	--	--------------	--------------------	--------------

<u>AB</u>		<u>300MG</u>	<u>A214387 003</u>	Feb 18, 2021
-----------	--	--------------	--------------------	--------------

<u>AB</u>	BIONPHARMA	<u>100MG</u>	<u>A213033 001</u>	Apr 28, 2021
-----------	------------	--------------	--------------------	--------------

<u>AB</u>		<u>200MG</u>	<u>A213033 002</u>	Apr 28, 2021
-----------	--	--------------	--------------------	--------------

<u>AB</u>		<u>300MG</u>	<u>A213033 003</u>	Apr 28, 2021
-----------	--	--------------	--------------------	--------------

<u>AB</u>	LUPIN PHARMS	<u>100MG</u>	<u>A211652 001</u>	Feb 18, 2021
-----------	--------------	--------------	--------------------	--------------

<u>AB</u>		<u>200MG</u>	<u>A211652 002</u>	Feb 18, 2021
-----------	--	--------------	--------------------	--------------

<u>AB</u>		<u>300MG</u>	<u>A211652 003</u>	Feb 18, 2021
-----------	--	--------------	--------------------	--------------

<u>AB</u>	MSN PHARMS INC	<u>100MG</u>	<u>A211741 001</u>	Feb 18, 2021
-----------	----------------	--------------	--------------------	--------------

<u>AB</u>		<u>200MG</u>	<u>A211741 002</u>	Feb 18, 2021
-----------	--	--------------	--------------------	--------------

<u>AB</u>		<u>300MG</u>	<u>A211741 003</u>	Feb 18, 2021
-----------	--	--------------	--------------------	--------------

<u>AB</u>	SCIEGEN PHARMS INC	<u>100MG</u>	<u>A214017 001</u>	Feb 18, 2021
-----------	--------------------	--------------	--------------------	--------------

<u>AB</u>		<u>200MG</u>	<u>A214017 002</u>	Feb 18, 2021
-----------	--	--------------	--------------------	--------------

<u>AB</u>		<u>300MG</u>	<u>A214017 003</u>	Feb 18, 2021
-----------	--	--------------	--------------------	--------------

<u>AB</u>	SOMERSET THERAPS LLC	<u>100MG</u>	<u>A214543 001</u>	May 05, 2021
-----------	----------------------	--------------	--------------------	--------------

<u>AB</u>		<u>200MG</u>	<u>A214543 002</u>	May 05, 2021
-----------	--	--------------	--------------------	--------------

<u>AB</u>		<u>300MG</u>	<u>A214543 003</u>	May 05, 2021
-----------	--	--------------	--------------------	--------------

<u>AB</u>	SUN PHARM	<u>100MG</u>	<u>A214384 001</u>	Feb 18, 2021
-----------	-----------	--------------	--------------------	--------------

<u>AB</u>		<u>200MG</u>	<u>A214384 002</u>	Feb 18, 2021
-----------	--	--------------	--------------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

DROXIDOPA

CAPSULE;ORAL

DROXIDOPA

<u>AB</u>		<u>300MG</u>	<u>A214384</u>	<u>003</u>	Feb 18, 2021
<u>AB</u>	ZYDUS PHARMS	<u>100MG</u>	<u>A211818</u>	<u>001</u>	Feb 18, 2021
<u>AB</u>		<u>200MG</u>	<u>A211818</u>	<u>002</u>	Feb 18, 2021
<u>AB</u>		<u>300MG</u>	<u>A211818</u>	<u>003</u>	Feb 18, 2021

NORTHERA

<u>AB</u>	+	LUNDBECK NA LTD	<u>100MG</u>	<u>N203202</u>	<u>001</u>	Feb 18, 2014
<u>AB</u>	+		<u>200MG</u>	<u>N203202</u>	<u>002</u>	Feb 18, 2014
<u>AB</u>	+		<u>300MG</u>	<u>N203202</u>	<u>003</u>	Feb 18, 2014

DULOXETINE HYDROCHLORIDE

CAPSULE, DELAYED REL PELLETS;ORAL

CYMBALTA

<u>AB</u>	+	LILLY	<u>EQ 20MG BASE</u>	<u>N021427</u>	<u>001</u>	Aug 03, 2004
<u>AB</u>	+		<u>EQ 30MG BASE</u>	<u>N021427</u>	<u>002</u>	Aug 03, 2004
<u>AB</u>	+		<u>EQ 60MG BASE</u>	<u>N021427</u>	<u>004</u>	Aug 03, 2004

DULOXETINE HYDROCHLORIDE

<u>AB</u>		ACTAVIS ELIZABETH	<u>EQ 20MG BASE</u>	<u>A090776</u>	<u>001</u>	Dec 17, 2013
<u>AB</u>			<u>EQ 30MG BASE</u>	<u>A090776</u>	<u>002</u>	Dec 17, 2013
<u>AB</u>			<u>EQ 60MG BASE</u>	<u>A090776</u>	<u>003</u>	Dec 17, 2013
<u>AB</u>		AJANTA PHARMA LTD	<u>EQ 20MG BASE</u>	<u>A208706</u>	<u>001</u>	Jan 06, 2017
<u>AB</u>			<u>EQ 30MG BASE</u>	<u>A208706</u>	<u>002</u>	Jan 06, 2017
<u>AB</u>			<u>EQ 40MG BASE</u>	<u>A208706</u>	<u>004</u>	Mar 11, 2019
<u>AB</u>			<u>EQ 60MG BASE</u>	<u>A208706</u>	<u>003</u>	Jan 06, 2017
<u>AB</u>		ALEMBIC	<u>EQ 20MG BASE</u>	<u>A202949</u>	<u>001</u>	Jun 09, 2014
<u>AB</u>			<u>EQ 30MG BASE</u>	<u>A202949</u>	<u>002</u>	Jun 09, 2014
<u>AB</u>			<u>EQ 60MG BASE</u>	<u>A202949</u>	<u>003</u>	Jun 09, 2014
<u>AB</u>		ALKEM LABS LTD	<u>EQ 20MG BASE</u>	<u>A203197</u>	<u>001</u>	Aug 26, 2015
<u>AB</u>			<u>EQ 30MG BASE</u>	<u>A203197</u>	<u>002</u>	Aug 26, 2015
<u>AB</u>			<u>EQ 60MG BASE</u>	<u>A203197</u>	<u>003</u>	Aug 26, 2015
<u>AB</u>		AUROBINDO PHARMA LTD	<u>EQ 20MG BASE</u>	<u>A090778</u>	<u>001</u>	Dec 11, 2013
<u>AB</u>			<u>EQ 30MG BASE</u>	<u>A090778</u>	<u>002</u>	Dec 11, 2013
<u>AB</u>			<u>EQ 60MG BASE</u>	<u>A090778</u>	<u>003</u>	Dec 11, 2013
<u>AB</u>		BRECKENRIDGE	<u>EQ 20MG BASE</u>	<u>A203088</u>	<u>001</u>	Jun 11, 2014
<u>AB</u>			<u>EQ 30MG BASE</u>	<u>A203088</u>	<u>002</u>	Jun 11, 2014
<u>AB</u>			<u>EQ 40MG BASE</u>	<u>A203088</u>	<u>004</u>	May 18, 2018
<u>AB</u>			<u>EQ 60MG BASE</u>	<u>A203088</u>	<u>003</u>	Jun 11, 2014
<u>AB</u>		CSPC OUYI	<u>EQ 20MG BASE</u>	<u>A211310</u>	<u>001</u>	Oct 16, 2018
<u>AB</u>			<u>EQ 30MG BASE</u>	<u>A211310</u>	<u>002</u>	Oct 16, 2018
<u>AB</u>			<u>EQ 60MG BASE</u>	<u>A211310</u>	<u>003</u>	Oct 16, 2018
<u>AB</u>		HETERO LABS LTD III	<u>EQ 20MG BASE</u>	<u>A204343</u>	<u>001</u>	Aug 03, 2016
<u>AB</u>			<u>EQ 30MG BASE</u>	<u>A204343</u>	<u>002</u>	Aug 03, 2016
<u>AB</u>			<u>EQ 60MG BASE</u>	<u>A204343</u>	<u>003</u>	Aug 03, 2016
<u>AB</u>		INVENTIA	<u>EQ 20MG BASE</u>	<u>A202336</u>	<u>001</u>	Oct 28, 2015
<u>AB</u>			<u>EQ 30MG BASE</u>	<u>A202336</u>	<u>002</u>	Oct 28, 2015
<u>AB</u>			<u>EQ 60MG BASE</u>	<u>A202336</u>	<u>003</u>	Oct 28, 2015
<u>AB</u>		LUPIN LTD	<u>EQ 20MG BASE</u>	<u>A090694</u>	<u>001</u>	Dec 11, 2013
<u>AB</u>			<u>EQ 30MG BASE</u>	<u>A090694</u>	<u>002</u>	Dec 11, 2013
<u>AB</u>			<u>EQ 40MG BASE</u>	<u>A090694</u>	<u>003</u>	Dec 11, 2013
<u>AB</u>			<u>EQ 60MG BASE</u>	<u>A090694</u>	<u>004</u>	Dec 11, 2013
<u>AB</u>		MACLEODS PHARMS LTD	<u>EQ 20MG BASE</u>	<u>A204815</u>	<u>001</u>	Mar 23, 2017
<u>AB</u>			<u>EQ 30MG BASE</u>	<u>A204815</u>	<u>002</u>	Mar 23, 2017
<u>AB</u>			<u>EQ 60MG BASE</u>	<u>A204815</u>	<u>003</u>	Mar 23, 2017
<u>AB</u>		PRINSTON INC	<u>EQ 20MG BASE</u>	<u>A206653</u>	<u>001</u>	May 18, 2017
<u>AB</u>			<u>EQ 30MG BASE</u>	<u>A206653</u>	<u>002</u>	May 18, 2017
<u>AB</u>			<u>EQ 60MG BASE</u>	<u>A206653</u>	<u>003</u>	May 18, 2017
<u>AB</u>		QINGDAO BAHEAL PHARM	<u>EQ 20MG BASE</u>	<u>A210599</u>	<u>001</u>	Apr 17, 2019
<u>AB</u>			<u>EQ 30MG BASE</u>	<u>A210599</u>	<u>002</u>	Apr 17, 2019
<u>AB</u>			<u>EQ 60MG BASE</u>	<u>A210599</u>	<u>003</u>	Apr 17, 2019
<u>AB</u>		SUNSHINE	<u>EQ 20MG BASE</u>	<u>A212328</u>	<u>001</u>	Feb 11, 2021
<u>AB</u>			<u>EQ 30MG BASE</u>	<u>A212328</u>	<u>002</u>	Feb 11, 2021
<u>AB</u>			<u>EQ 60MG BASE</u>	<u>A212328</u>	<u>003</u>	Feb 11, 2021
<u>AB</u>		TORRENT	<u>EQ 20MG BASE</u>	<u>A090774</u>	<u>001</u>	Dec 11, 2013
<u>AB</u>			<u>EQ 30MG BASE</u>	<u>A090774</u>	<u>002</u>	Dec 11, 2013
<u>AB</u>			<u>EQ 60MG BASE</u>	<u>A090774</u>	<u>003</u>	Dec 11, 2013
<u>AB</u>		ZYDUS HLTHCARE	<u>EQ 20MG BASE</u>	<u>A090739</u>	<u>001</u>	Jan 08, 2014
<u>AB</u>			<u>EQ 30MG BASE</u>	<u>A090739</u>	<u>002</u>	Jan 08, 2014
<u>AB</u>			<u>EQ 40MG BASE</u>	<u>A090739</u>	<u>004</u>	Apr 18, 2023
<u>AB</u>			<u>EQ 60MG BASE</u>	<u>A090739</u>	<u>003</u>	Jan 08, 2014

PRESCRIPTION DRUG PRODUCT LIST

DULOXETINE HYDROCHLORIDE

CAPSULE, DELAYED REL PELLETS;ORAL

DULOXETINE HYDROCHLORIDE

AB	ZYDUS PHARMS	EQ 20MG BASE	A090728 001	Jan 08, 2014
AB		EQ 30MG BASE	A090728 002	Jan 08, 2014
AB		EQ 60MG BASE	A090728 003	Jan 08, 2014
	DRIZALMA SPRINKLE			
	+ SUN PHARM	EQ 20MG BASE	N212516 001	Jul 19, 2019
	+	EQ 30MG BASE	N212516 002	Jul 19, 2019
	+	EQ 40MG BASE	N212516 003	Jul 19, 2019
	+!	EQ 60MG BASE	N212516 004	Jul 19, 2019

DURLOBACTAM SODIUM; DURLOBACTAM SODIUM; SULBACTAM SODIUM

POWDER; INTRAVENOUS

XACDURO (COPACKAGED)

	+! ENTASIS THERAP	EQ 500MG BASE/VIAL; EQ 500MG BASE/VIAL;EQ 1GM BASE/VIAL	N216974 001	May 23, 2023
--	-------------------	--	-------------	--------------

DUTASTERIDE

CAPSULE;ORAL

AVODART

AB	+! WAYLIS THERAP	0.5MG	N021319 001	Nov 20, 2001
-----------	------------------	--------------	--------------------	--------------

DUTASTERIDE

AB	ACELLA	0.5MG	A206373 001	Mar 17, 2016
AB	ADAPTIS	0.5MG	A204376 001	Apr 07, 2017
AB	AMNEAL PHARMS	0.5MG	A203118 001	Nov 20, 2015
AB	ASCENT PHARMS INC	0.5MG	A206574 001	Oct 21, 2016
AB	AUROBINDO PHARMA LTD	0.5MG	A202660 001	Nov 20, 2015
AB	BARR	0.5MG	A090095 001	Dec 21, 2010
AB	CHARTWELL	0.5MG	A200899 001	Nov 20, 2015
AB	HUMANWELL PURACAP	0.5MG	A209909 001	Nov 21, 2017
AB	STRIDES SOFTGELS	0.5MG	A204262 001	Nov 20, 2015
AB	ZYDUS LIFESCIENCES	0.5MG	A204373 001	Oct 04, 2017

DUTASTERIDE; TAMSULOSIN HYDROCHLORIDE

CAPSULE;ORAL

DUTASTERIDE AND TAMSULOSIN HYDROCHLORIDE

AB	AUROBINDO PHARMA	0.5MG;0.4MG	A213300 001	Jul 18, 2024
AB	ENDO OPERATIONS	0.5MG;0.4MG	A202509 001	Feb 26, 2014
AB	ZYDUS PHARMS	0.5MG;0.4MG	A207769 001	May 24, 2018
	JALYN			
AB	+! WAYLIS THERAP	0.5MG;0.4MG	N022460 001	Jun 14, 2010

DUVELISIB

CAPSULE;ORAL

COPIKTRA

	+ SECURA	15MG	N211155 001	Sep 24, 2018
	+	25MG	N211155 002	Sep 24, 2018

DYCLONINE HYDROCHLORIDE

SOLUTION;TOPICAL

DYCLOPRO

	! SEPTODONT	0.5%	A200480 001	Nov 20, 2018
	!	1%	A200480 002	Nov 20, 2018

ECHOTHIOPHATE IODIDE

FOR SOLUTION;OPHTHALMIC

PHOSPHOLINE IODIDE

	+! FERA PHARMS LLC	0.125%	N011963 001	
--	--------------------	--------	-------------	--

ECONAZOLE NITRATE

AEROSOL, FOAM;TOPICAL

ECOZA

	+! RESILIA PHARMS	1%	N205175 001	Oct 24, 2013
--	-------------------	----	-------------	--------------

CREAM;TOPICAL

ECONAZOLE NITRATE

AB	! PADAGIS ISRAEL	1%	A076479 001	Jun 23, 2004
AB	TARO	1%	A076005 001	Nov 26, 2002

EDARAVONE

SOLUTION;INTRAVENOUS

EDARAVONE

AP	DR REDDYS	60MG/100ML (0.6MG/ML)	A215917 001	May 06, 2024
AP	GLAND PHARMA LTD	30MG/100ML (0.3MG/ML)	A216199 001	May 06, 2024
AP		60MG/100ML (0.6MG/ML)	A216199 002	May 06, 2024
AP	HIKMA	30MG/100ML (0.3MG/ML)	A215508 001	May 06, 2024

PRESCRIPTION DRUG PRODUCT LIST

EDARAVONE

SOLUTION; INTRAVENOUS

EDARAVONE

AP	LONG GROVE PHARMS	<u>30MG/100ML (0.3MG/ML)</u>	A218354 001	May 06, 2024
AP	SANDOZ	<u>30MG/100ML (0.3MG/ML)</u>	A216902 001	Dec 23, 2024
AP	XGEN PHARMS	<u>30MG/100ML (0.3MG/ML)</u>	A217565 001	Oct 31, 2024
AP		<u>60MG/100ML (0.6MG/ML)</u>	A217565 002	Oct 31, 2024

RADICAVA

AP	+!	! MITSUBISHI TANABE	<u>30MG/100ML (0.3MG/ML)</u>	N209176 001	May 05, 2017
AP	+!		<u>60MG/100ML (0.6MG/ML)</u>	N209176 002	Nov 15, 2018

SUSPENSION; ORAL

RADICAVA ORS

	+!	! MITSUBISHI TANABE	105MG/5ML	N215446 001	May 12, 2022
--	----	---------------------	-----------	-------------	--------------

EDETATE CALCIUM DISODIUM

INJECTABLE; INJECTION

EDETATE CALCIUM DISODIUM

	!	CASPER PHARMA LLC	200MG/ML	A216435 001	May 03, 2023
--	---	-------------------	----------	-------------	--------------

EDOXYBAN TOSYLATE

TABLET; ORAL

SAVAYSA

	+	DAIICHI SANKYO INC	EQ 15MG BASE	N206316 001	Jan 08, 2015
	+		EQ 30MG BASE	N206316 002	Jan 08, 2015
	+!		EQ 60MG BASE	N206316 003	Jan 08, 2015

EFAVIRENZ

CAPSULE; ORAL

EFAVIRENZ

		AUROBINDO PHARMA	50MG	A078064 001	Dec 15, 2017
			100MG	A078064 002	Dec 15, 2017
	!		200MG	A078064 003	Dec 15, 2017

TABLET; ORAL

EFAVIRENZ

AB		AUROBINDO PHARMA LTD	600MG	A077673 001	Sep 21, 2018
AB		CIPLA	600MG	A204766 001	Jun 15, 2018
AB	!	HETERO LABS LTD III	600MG	A078886 001	Apr 27, 2018
AB		MACLEODS PHARMS LTD	600MG	A091579 001	May 10, 2024
AB		STRIDES PHARMA	600MG	A204869 001	Mar 12, 2018

EFAVIRENZ; EMTRICITABINE; TENOFOVIR DISOPROXIL FUMARATE

TABLET; ORAL

EFAVIRENZ, EMTRICITABINE, AND TENOFOVIR DISOPROXIL FUMARATE

AB	!	AUROBINDO PHARMA	<u>600MG; 200MG; 300MG</u>	A203041 001	Sep 04, 2018
AB		CIPLA	<u>600MG; 200MG; 300MG</u>	A206894 001	Jun 03, 2019
AB		HETERO LABS LTD V	<u>600MG; 200MG; 300MG</u>	A203053 001	Jan 24, 2022
AB		LAURUS	<u>600MG; 200MG; 300MG</u>	A213541 001	Dec 22, 2021
AB		MACLEODS PHARMS LTD	<u>600MG; 200MG; 300MG</u>	A204287 001	Sep 13, 2021
AB		TEVA PHARMS USA	<u>600MG; 200MG; 300MG</u>	A091215 001	Nov 09, 2018

EFAVIRENZ; LAMIVUDINE; TENOFOVIR DISOPROXIL FUMARATE

TABLET; ORAL

EFAVIRENZ, LAMIVUDINE AND TENOFOVIR DISOPROXIL FUMARATE

AB		LAURUS	<u>400MG; 300MG; 300MG</u>	A213038 001	May 14, 2020
AB			<u>600MG; 300MG; 300MG</u>	A212786 001	May 14, 2020
		SYMFI			
AB	+!	MYLAN LABS LTD	<u>600MG; 300MG; 300MG</u>	N022142 001	Mar 22, 2018
		SYMFI LO			
AB	+!	MYLAN	<u>400MG; 300MG; 300MG</u>	N208255 001	Feb 05, 2018

EFINACONAZOLE

SOLUTION; TOPICAL

EFINACONAZOLE

AB		TEVA PHARMS USA	10%	A211827 001	Dec 16, 2020
AB		UMEDICA	10%	A211969 001	Jun 21, 2021

JUBLIA

AB	+!	BAUSCH	10%	N203567 001	Jun 06, 2014
-----------	----	--------	------------	--------------------	--------------

EFLORNITHINE HYDROCHLORIDE

TABLET; ORAL

IWILFIN

	+!	USWM	EQ 192MG BASE	N215500 001	Dec 13, 2023
--	----	------	---------------	-------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

ELACESTRANT HYDROCHLORIDE

TABLET; ORAL

ORSERDU

+	STEMLINE THERAP	EQ 86MG BASE	N217639 001	Jan 27, 2023
+	!	EQ 345MG BASE	N217639 002	Jan 27, 2023

ELAFIBRANOR

TABLET; ORAL

IQIRVO

+	IPSEN	80MG	N218860 001	Jun 10, 2024
---	-------	------	-------------	--------------

ELAGOLIX SODIUM

TABLET; ORAL

ORILISSA

+	ABBVIE	EQ 150MG BASE	N210450 001	Jul 23, 2018
+	!	EQ 200MG BASE	N210450 002	Jul 23, 2018

ELAGOLIX SODIUM, ESTRADIOL, NORETHINDRONE ACETATE; ELAGOLIX SODIUM

CAPSULE; ORAL

ORIAHNN (COPACKAGED)

+	ABBVIE	EQ 300MG BASE, 1MG, 0.5MG; EQ 300MG BASE	N213388 001	May 29, 2020
---	--------	--	-------------	--------------

ELBASVIR; GRAZOPREVIR

TABLET; ORAL

ZEPATIER

+	MSD SUB MERCK	50MG; 100MG	N208261 001	Jan 28, 2016
---	---------------	-------------	-------------	--------------

ELETRIPTAN HYDROBROMIDE

TABLET; ORAL

ELETRIPTAN HYDROBROMIDE

<u>AB</u>	AUROBINDO PHARMA	<u>EQ 20MG BASE</u>	<u>A210708 001</u>	Jan 15, 2019
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A210708 002</u>	Jan 15, 2019
<u>AB</u>	BEXIMCO PHARMS USA	<u>EQ 20MG BASE</u>	<u>A215467 001</u>	Jul 13, 2022
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A215467 002</u>	Jul 13, 2022
<u>AB</u>	CHARTWELL RX	<u>EQ 20MG BASE</u>	<u>A205186 001</u>	Aug 29, 2017
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A205186 002</u>	Aug 29, 2017
<u>AB</u>	MYLAN	<u>EQ 20MG BASE</u>	<u>A205152 001</u>	Aug 11, 2017
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A205152 002</u>	Aug 11, 2017
<u>AB</u>	TEVA PHARMS USA	<u>EQ 20MG BASE</u>	<u>A202040 001</u>	Jun 27, 2017
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A202040 002</u>	Jun 27, 2017
<u>AB</u>	ZYDUS PHARMS	<u>EQ 20MG BASE</u>	<u>A206409 001</u>	Jun 16, 2017
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A206409 002</u>	Jun 16, 2017

RELPAK

<u>AB</u>	+	UPJOHN	<u>EQ 20MG BASE</u>	<u>N021016 001</u>	Dec 26, 2002
<u>AB</u>	+	!	<u>EQ 40MG BASE</u>	<u>N021016 002</u>	Dec 26, 2002

ELEXACAFTOR, IVACAFTOR, TEZACAFTOR; IVACAFTOR

GRANULES; ORAL

TRIKAFTA (COPACKAGED)

+	VERTEX PHARMS INC	80MG, 60MG, 40MG; 59.5MG	N217660 001	Apr 26, 2023
+	!	100MG, 75MG, 50MG; 75MG	N217660 002	Apr 26, 2023

TABLET; ORAL

TRIKAFTA (COPACKAGED)

+	VERTEX PHARMS INC	50MG, 37.5MG, 25MG; 75MG	N212273 002	Jun 08, 2021
+	!	100MG, 75MG, 50MG; 150MG	N212273 001	Oct 21, 2019

ELIGLUSTAT TARTRATE

CAPSULE; ORAL

CERDELGA

<u>AB</u>	+	GENZYME CORP	<u>EQ 84MG BASE</u>	<u>N205494 001</u>	Aug 19, 2014
------------------	---	--------------	----------------------------	---------------------------	--------------

ELIGLUSTAT TARTRATE

<u>AB</u>	AIZANT	<u>EQ 84MG BASE</u>	<u>A212463 001</u>	Sep 08, 2021
<u>AB</u>	APOTEX	<u>EQ 84MG BASE</u>	<u>A212425 001</u>	Jul 10, 2024
<u>AB</u>	CIPLA	<u>EQ 84MG BASE</u>	<u>A212369 001</u>	Dec 10, 2024
<u>AB</u>	UPSHER SMITH LABS	<u>EQ 84MG BASE</u>	<u>A212420 001</u>	May 02, 2023

ELTROMBOPAG CHOLINE

TABLET; ORAL

ALVAIZ

+	TEVA PHARMS INC	EQ 9MG BASE	N216774 001	Nov 29, 2023
+		EQ 18MG BASE	N216774 002	Nov 29, 2023
+		EQ 36MG BASE	N216774 003	Nov 29, 2023
+	!	EQ 54MG BASE	N216774 004	Nov 29, 2023

PRESCRIPTION DRUG PRODUCT LIST

ELTROMBOPAG OLAMINE

FOR SUSPENSION;ORAL

ELTROMBOPAG OLAMINE

AB	ANNORA PHARMA	EQ 12.5MG ACID/PACKET	A216620 001	Apr 18, 2024
AB		EQ 25MG ACID/PACKET	A216620 002	Apr 18, 2024

PROMACTA KIT

AB	+ NOVARTIS	EQ 12.5MG ACID/PACKET	N207027 002	Sep 27, 2018
AB	+!	EQ 25MG ACID/PACKET	N207027 001	Aug 24, 2015

TABLET;ORAL

PROMACTA

+	NOVARTIS	EQ 12.5MG ACID	N022291 004	Oct 20, 2011
+		EQ 25MG ACID	N022291 001	Nov 20, 2008
+		EQ 50MG ACID	N022291 002	Nov 20, 2008
+	!	EQ 75MG ACID	N022291 003	Sep 08, 2009

ELUXADOLINE

TABLET;ORAL

VIBERZI

+	ABBVIE	75MG	N206940 001	May 27, 2015
+	!	100MG	N206940 002	May 27, 2015

EMPAGLIFLOZIN

TABLET;ORAL

JARDIANCE

+	BOEHRINGER INGELHEIM	10MG	N204629 001	Aug 01, 2014
+	!	25MG	N204629 002	Aug 01, 2014

EMPAGLIFLOZIN; LINAGLIPTIN

TABLET;ORAL

GLYXAMBI

+	BOEHRINGER INGELHEIM	10MG;5MG	N206073 001	Jan 30, 2015
+	!	25MG;5MG	N206073 002	Jan 30, 2015

EMPAGLIFLOZIN; LINAGLIPTIN; METFORMIN HYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL

TRIJARDY XR

+	BOEHRINGER INGELHEIM	5MG;2.5MG;1GM	N212614 001	Jan 27, 2020
+		10MG;5MG;1GM	N212614 002	Jan 27, 2020
+		12.5MG;2.5MG;1GM	N212614 003	Jan 27, 2020
+	!	25MG;5MG;1GM	N212614 004	Jan 27, 2020

EMPAGLIFLOZIN; METFORMIN HYDROCHLORIDE

TABLET;ORAL

SYNJARDY

+	BOEHRINGER INGELHEIM	5MG;500MG	N206111 001	Aug 26, 2015
+		5MG;1GM	N206111 002	Aug 26, 2015
+		12.5MG;500MG	N206111 003	Aug 26, 2015
+	!	12.5MG;1GM	N206111 004	Aug 26, 2015

TABLET, EXTENDED RELEASE;ORAL

SYNJARDY XR

+	BOEHRINGER INGELHEIM	5MG;1GM	N208658 001	Dec 09, 2016
+		10MG;1GM	N208658 002	Dec 09, 2016
+		12.5MG;1GM	N208658 003	Dec 09, 2016
+	!	25MG;1GM	N208658 004	Dec 09, 2016

EMTRICITABINE

CAPSULE;ORAL

EMTRICITABINE

AB	AUROBINDO PHARMA LTD	200MG	A079188 001	Mar 15, 2023
-----------	-------------------------	--------------	--------------------	--------------

AB	CIPLA	200MG	A091168 001	Jul 02, 2018
-----------	-------	--------------	--------------------	--------------

EMTRIVA

AB	+! GILEAD	200MG	N021500 001	Jul 02, 2003
-----------	-----------	--------------	--------------------	--------------

SOLUTION;ORAL

EMTRIVA

+	GILEAD	10MG/ML	N021896 001	Sep 28, 2005
---	--------	---------	-------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

EMTRICITABINE; RILPIVIRINE HYDROCHLORIDE; TENOFOVIR ALAFENAMIDE FUMARATE

TABLET;ORAL

ODEFSEY

+! GILEAD SCIENCES INC 200MG;EQ 25MG BASE;EQ 25MG BASE N208351 001 Mar 01, 2016

EMTRICITABINE; RILPIVIRINE HYDROCHLORIDE; TENOFOVIR DISOPROXIL FUMARATE

TABLET;ORAL

COMPLERA

+! GILEAD SCIENCES INC 200MG;EQ 25MG BASE;300MG N202123 001 Aug 10, 2011

EMTRICITABINE; TENOFOVIR ALAFENAMIDE FUMARATE

TABLET;ORAL

DESCOVY**AB** +! GILEAD SCIENCES INC 200MG;EQ 25MG BASE **N208215 001** Apr 04, 2016EMTRICITABINE AND TENOFOVIR ALAFENAMIDE FUMARATE**AB** LUPIN LTD 200MG;EQ 25MG BASE **A213926 001** Dec 13, 2024

DESCOVY

+ GILEAD SCIENCES INC 120MG;EQ 15MG BASE N208215 002 Jan 07, 2022

EMTRICITABINE; TENOFOVIR DISOPROXIL FUMARATE

TABLET;ORAL

EMTRICITABINE AND TENOFOVIR DISOPROXIL FUMARATE**AB** HETERO LABS LTD III 200MG;300MG **A201806 001** Oct 07, 2021EMTRICITABINE AND TENOFOVIR DISOPROXIL FUMARATE**AB** AMNEAL PHARMS CO 100MG;150MG **A209721 001** Aug 22, 2018**AB** 133MG;200MG **A209721 002** Aug 22, 2018**AB** 167MG;250MG **A209721 003** Aug 22, 2018**AB** 200MG;300MG **A209721 004** Aug 22, 2018**AB** AUROBINDO PHARMA 200MG;300MG **A090513 001** Jan 26, 2018**AB** AUROBINDO PHARMA 100MG;150MG **A211640 001** Mar 09, 2023

LTD

AB 133MG;200MG **A211640 002** Mar 09, 2023**AB** 167MG;250MG **A211640 003** Mar 09, 2023**AB** CHARTWELL RX 200MG;300MG **A204131 001** Jun 04, 2021**AB** LAURUS 200MG;300MG **A212114 001** Jul 26, 2019**AB** MACLEODS PHARMS LTD 200MG;300MG **A203442 001** May 15, 2020**AB** MYLAN 200MG;300MG **A206436 001** Apr 09, 2018**AB** STRIDES PHARMA 200MG;300MG **A091055 001** Jan 13, 2021**AB** TEVA PHARMS USA 200MG;300MG **A090894 001** Jun 08, 2017**AB** ZYDUS PHARMS 100MG;150MG **A212689 002** Jul 01, 2021**AB** 133MG;200MG **A212689 003** Jul 01, 2021**AB** 167MG;250MG **A212689 004** Jul 01, 2021**AB** 200MG;300MG **A212689 001** Feb 28, 2020TRUVADA**AB** + GILEAD 100MG;150MG **N021752 002** Mar 10, 2016**AB** + 133MG;200MG **N021752 003** Mar 10, 2016**AB** + 167MG;250MG **N021752 004** Mar 10, 2016**AB** +! 200MG;300MG **N021752 001** Aug 02, 2004ENALAPRIL MALEATE

SOLUTION;ORAL

ENALAPRIL MALEATE**AB** ALKEM LABS LTD 1MG/ML **A213714 001** Mar 30, 2022**AB** AMNEAL 1MG/ML **A212894 001** Jun 29, 2022**AB** ANNORA PHARMA 1MG/ML **A214467 001** Feb 24, 2022**AB** AUROBINDO PHARMA 1MG/ML **A216458 001** Jan 08, 2024**AB** BIONPHARMA 1MG/ML **A212408 001** Aug 10, 2021EPANED**AB** +! AZURITY 1MG/ML **N208686 001** Sep 20, 2016

TABLET;ORAL

ENALAPRIL MALEATE**AB** AIPING PHARM INC 2.5MG **A075178 002** Mar 23, 2001**AB** 5MG **A075178 001** Mar 23, 2001**AB** 10MG **A075178 003** Mar 23, 2001**AB** 20MG **A075178 004** Mar 23, 2001**AB** HERITAGE PHARMA 2.5MG **A075479 001** Aug 22, 2000**AB** 5MG **A075479 002** Aug 22, 2000**AB** 10MG **A075479 003** Aug 22, 2000**AB** 20MG **A075479 004** Aug 22, 2000**AB** PRINSTON INC 2.5MG **A213273 001** Jul 07, 2022**AB** 5MG **A213273 002** Jul 07, 2022**AB** 10MG **A213273 003** Jul 07, 2022**AB** 20MG **A213273 004** Jul 07, 2022**AB** SANDOZ INC 2.5MG **A075496 001** Aug 22, 2000**AB** 5MG **A075496 002** Aug 22, 2000

PRESCRIPTION DRUG PRODUCT LIST

ENALAPRIL MALEATE

TABLET; ORAL

ENALAPRIL MALEATE

<u>AB</u>		<u>10MG</u>	<u>A075459 001</u>	Aug 22, 2000
<u>AB</u>		<u>20MG</u>	<u>A075459 002</u>	Aug 22, 2000
<u>AB</u>	TARO	<u>2.5MG</u>	<u>A075657 001</u>	Jan 23, 2001
<u>AB</u>		<u>5MG</u>	<u>A075657 002</u>	Jan 23, 2001
<u>AB</u>		<u>10MG</u>	<u>A075657 003</u>	Jan 23, 2001
<u>AB</u>		<u>20MG</u>	<u>A075657 004</u>	Jan 23, 2001
<u>AB</u>	UNIQUE PHARM	<u>2.5MG</u>	<u>A218531 001</u>	Sep 19, 2024
<u>AB</u>		<u>5MG</u>	<u>A218531 002</u>	Sep 19, 2024
<u>AB</u>		<u>10MG</u>	<u>A218531 003</u>	Sep 19, 2024
<u>AB</u>		<u>20MG</u>	<u>A218531 004</u>	Sep 19, 2024
<u>AB</u>	WOCKHARDT LTD	<u>2.5MG</u>	<u>A075483 001</u>	Aug 22, 2000
<u>AB</u>		<u>5MG</u>	<u>A075483 002</u>	Aug 22, 2000
<u>AB</u>		<u>10MG</u>	<u>A075483 003</u>	Aug 22, 2000
<u>AB</u>		<u>20MG</u>	<u>A075483 004</u>	Aug 22, 2000

VASOTEC

<u>AB</u>	+	BAUSCH	<u>2.5MG</u>	<u>N018998 005</u>	Jul 26, 1988
<u>AB</u>	+		<u>5MG</u>	<u>N018998 001</u>	Dec 24, 1985
<u>AB</u>	+		<u>10MG</u>	<u>N018998 002</u>	Dec 24, 1985
<u>AB</u>	+	!	<u>20MG</u>	<u>N018998 003</u>	Dec 24, 1985

ENALAPRIL MALEATE; HYDROCHLOROTHIAZIDE

TABLET; ORAL

ENALAPRIL MALEATE AND HYDROCHLOROTHIAZIDE

<u>AB</u>	DR REDDYS LABS LTD	<u>5MG;12.5MG</u>	<u>A075909 001</u>	Oct 15, 2001
<u>AB</u>		<u>10MG;25MG</u>	<u>A075909 002</u>	Oct 15, 2001
<u>AB</u>	TARO PHARM INDS	<u>5MG;12.5MG</u>	<u>A075788 001</u>	Sep 18, 2001
<u>AB</u>		<u>10MG;25MG</u>	<u>A075788 002</u>	Sep 18, 2001

VASERETIC

<u>AB</u>	+	BAUSCH	<u>5MG;12.5MG</u>	<u>N019221 003</u>	Jul 12, 1995
<u>AB</u>	+	!	<u>10MG;25MG</u>	<u>N019221 001</u>	Oct 31, 1986

ENALAPRILAT

INJECTABLE; INJECTION

ENALAPRILAT

<u>AP</u>	CHARTWELL	<u>1.25MG/ML</u>	<u>A075634 001</u>	Aug 22, 2000	
	INJECTABLE				
<u>AP</u>	DR REDDYS	<u>1.25MG/ML</u>	<u>A075578 001</u>	Aug 22, 2000	
<u>AP</u>	!	HIKMA FARMACEUTICA	<u>1.25MG/ML</u>	<u>A078687 001</u>	Dec 23, 2008
<u>AP</u>	HOSPIRA	<u>1.25MG/ML</u>	<u>A075458 001</u>	Aug 22, 2000	

ENASIDENIB MESYLATE

TABLET; ORAL

IDHIFA

+	BRISTOL MYERS	EQ 50MG BASE	N209606 001	Aug 01, 2017
	SQUIBB			
+	!	EQ 100MG BASE	N209606 002	Aug 01, 2017

ENCORAFENIB

CAPSULE; ORAL

BRAFTOVI

+	ARRAY BIOPHARMA INC	75MG	N210496 002	Jun 27, 2018
---	---------------------	------	-------------	--------------

ENFUVRTIDE

INJECTABLE; SUBCUTANEOUS

FUZEON

+	ROCHE	90MG/VIAL	N021481 001	Mar 13, 2003
---	-------	-----------	-------------	--------------

ENOXAPARIN SODIUM

INJECTABLE; INTRAVENOUS, SUBCUTANEOUS

ENOXAPARIN SODIUM

<u>AB</u>	AMPHASTAR PHARMS	<u>300MG/3ML (100MG/ML)</u>	<u>A208600 001</u>	Mar 14, 2019
	INC			
<u>AB</u>	NANJING KING-FRIEND	<u>300MG/3ML (100MG/ML)</u>	<u>A214856 001</u>	Jun 14, 2022
<u>AB</u>	SANDOZ INC	<u>300MG/3ML (100MG/ML)</u>	<u>A078660 001</u>	Nov 28, 2011

LOVENOX

<u>AB</u>	+	SANOFI AVENTIS US	<u>300MG/3ML (100MG/ML)</u>	<u>N020164 009</u>	Jan 23, 2003
-----------	---	-------------------	-----------------------------	--------------------	--------------

INJECTABLE; SUBCUTANEOUS

ENOXAPARIN SODIUM (PRESERVATIVE FREE)

<u>AP</u>	AMPHASTAR PHARM	<u>30MG/0.3ML (100MG/ML)</u>	<u>A076684 001</u>	Sep 19, 2011
<u>AP</u>		<u>40MG/0.4ML (100MG/ML)</u>	<u>A076684 002</u>	Sep 19, 2011
<u>AP</u>		<u>60MG/0.6ML (100MG/ML)</u>	<u>A076684 003</u>	Sep 19, 2011
<u>AP</u>		<u>80MG/0.8ML (100MG/ML)</u>	<u>A076684 004</u>	Sep 19, 2011
<u>AP</u>		<u>100MG/ML (100MG/ML)</u>	<u>A076684 005</u>	Sep 19, 2011

PRESCRIPTION DRUG PRODUCT LIST

ENOXAPARIN SODIUM

INJECTABLE; SUBCUTANEOUS

ENOXAPARIN SODIUM (PRESERVATIVE FREE)

<u>AP</u>		<u>120MG/0.8ML (150MG/ML)</u>	<u>A076684 006</u>	Sep 19, 2011
<u>AP</u>		<u>150MG/ML (150MG/ML)</u>	<u>A076684 007</u>	Sep 19, 2011
<u>AP</u>	BE PHARMS	<u>30MG/0.3ML (100MG/ML)</u>	<u>A214646 001</u>	Jun 06, 2023
<u>AP</u>		<u>40MG/0.4ML (100MG/ML)</u>	<u>A214646 002</u>	Jun 06, 2023
<u>AP</u>		<u>60MG/0.6ML (100MG/ML)</u>	<u>A214646 003</u>	Jun 06, 2023
<u>AP</u>		<u>80MG/0.8ML (100MG/ML)</u>	<u>A214646 004</u>	Jun 06, 2023
<u>AP</u>		<u>100MG/ML (100MG/ML)</u>	<u>A214646 005</u>	Jun 06, 2023
<u>AP</u>		<u>120MG/0.8ML (150MG/ML)</u>	<u>A214646 006</u>	Jun 06, 2023
<u>AP</u>		<u>150MG/ML (150MG/ML)</u>	<u>A214646 007</u>	Jun 06, 2023
<u>AP</u>	GLAND	<u>30MG/0.3ML (100MG/ML)</u>	<u>A078990 001</u>	Sep 28, 2018
<u>AP</u>		<u>40MG/0.4ML (100MG/ML)</u>	<u>A078990 002</u>	Sep 28, 2018
<u>AP</u>		<u>60MG/0.6ML (100MG/ML)</u>	<u>A078990 003</u>	Sep 28, 2018
<u>AP</u>		<u>80MG/0.8ML (100MG/ML)</u>	<u>A078990 004</u>	Sep 28, 2018
<u>AP</u>		<u>100MG/ML (100MG/ML)</u>	<u>A078990 005</u>	Sep 28, 2018
<u>AP</u>		<u>120MG/0.8ML (150MG/ML)</u>	<u>A078990 006</u>	Sep 28, 2018
<u>AP</u>		<u>150MG/ML (150MG/ML)</u>	<u>A078990 007</u>	Sep 28, 2018
<u>AP</u>	NANJING KING-FRIEND	<u>30MG/0.3ML (100MG/ML)</u>	<u>A206834 001</u>	Nov 29, 2019
<u>AP</u>		<u>40MG/0.4ML (100MG/ML)</u>	<u>A206834 002</u>	Nov 29, 2019
<u>AP</u>		<u>60MG/0.6ML (100MG/ML)</u>	<u>A206834 003</u>	Nov 29, 2019
<u>AP</u>		<u>80MG/0.8ML (100MG/ML)</u>	<u>A206834 004</u>	Nov 29, 2019
<u>AP</u>		<u>100MG/ML (100MG/ML)</u>	<u>A206834 005</u>	Nov 29, 2019
<u>AP</u>		<u>120MG/0.8ML (150MG/ML)</u>	<u>A206834 006</u>	Nov 29, 2019
<u>AP</u>		<u>150MG/ML (150MG/ML)</u>	<u>A206834 007</u>	Nov 29, 2019
<u>AP</u>	SANDOZ	<u>30MG/0.3ML (100MG/ML)</u>	<u>A077857 002</u>	Jul 23, 2010
<u>AP</u>		<u>40MG/0.4ML (100MG/ML)</u>	<u>A077857 003</u>	Jul 23, 2010
<u>AP</u>		<u>60MG/0.6ML (100MG/ML)</u>	<u>A077857 004</u>	Jul 23, 2010
<u>AP</u>		<u>80MG/0.8ML (100MG/ML)</u>	<u>A077857 005</u>	Jul 23, 2010
<u>AP</u>		<u>100MG/ML (100MG/ML)</u>	<u>A077857 001</u>	Jul 23, 2010
<u>AP</u>		<u>120MG/0.8ML (150MG/ML)</u>	<u>A077857 006</u>	Jul 23, 2010
<u>AP</u>		<u>150MG/ML (150MG/ML)</u>	<u>A077857 007</u>	Jul 23, 2010
<u>AP</u>	SHENZHEN TECHDOW	<u>30MG/0.3ML (100MG/ML)</u>	<u>A205660 001</u>	Mar 15, 2023
<u>AP</u>		<u>40MG/0.4ML (100MG/ML)</u>	<u>A205660 002</u>	Mar 15, 2023
<u>AP</u>		<u>60MG/0.6ML (100MG/ML)</u>	<u>A205660 003</u>	Mar 15, 2023
<u>AP</u>		<u>80MG/0.8ML (100MG/ML)</u>	<u>A205660 004</u>	Mar 15, 2023
<u>AP</u>		<u>100MG/ML (100MG/ML)</u>	<u>A205660 005</u>	Mar 15, 2023
<u>AP</u>		<u>120MG/0.8ML (150MG/ML)</u>	<u>A205660 006</u>	Mar 15, 2023
<u>AP</u>		<u>150MG/ML (150MG/ML)</u>	<u>A205660 007</u>	Mar 15, 2023
<u>AP</u>	ZYDUS PHARMS	<u>30MG/0.3ML (100MG/ML)</u>	<u>A076726 001</u>	Jun 23, 2014
<u>AP</u>		<u>40MG/0.4ML (100MG/ML)</u>	<u>A076726 002</u>	Jun 23, 2014
<u>AP</u>		<u>60MG/0.6ML (100MG/ML)</u>	<u>A076726 003</u>	Jun 23, 2014
<u>AP</u>		<u>80MG/0.8ML (100MG/ML)</u>	<u>A076726 004</u>	Jun 23, 2014
<u>AP</u>		<u>100MG/ML (100MG/ML)</u>	<u>A076726 005</u>	Jun 23, 2014
<u>AP</u>		<u>120MG/0.8ML (150MG/ML)</u>	<u>A076726 006</u>	Jun 23, 2014
<u>AP</u>		<u>150MG/ML (150MG/ML)</u>	<u>A076726 007</u>	Jun 23, 2014

LOVENOX (PRESERVATIVE FREE)

<u>AP</u>	+	SANOFI AVENTIS US	<u>30MG/0.3ML (100MG/ML)</u>	<u>N020164 001</u>	Mar 29, 1993
<u>AP</u>	+		<u>40MG/0.4ML (100MG/ML)</u>	<u>N020164 002</u>	Jan 30, 1998
<u>AP</u>	+		<u>60MG/0.6ML (100MG/ML)</u>	<u>N020164 003</u>	Mar 27, 1998
<u>AP</u>	+		<u>80MG/0.8ML (100MG/ML)</u>	<u>N020164 004</u>	Mar 27, 1998
<u>AP</u>	+		<u>100MG/ML (100MG/ML)</u>	<u>N020164 005</u>	Mar 27, 1998
<u>AP</u>	+		<u>120MG/0.8ML (150MG/ML)</u>	<u>N020164 007</u>	Jun 02, 2000
<u>AP</u>	+		<u>150MG/ML (150MG/ML)</u>	<u>N020164 008</u>	Jun 02, 2000

ENSARTINIB HYDROCHLORIDE

CAPSULE; ORAL

ENSACOVE

+ XCOVERY

EQ 25MG BASE

N218171 001 Dec 18, 2024

+!

EQ 100MG BASE

N218171 002 Dec 18, 2024

ENSIFENTRINE

SUSPENSION; INHALATION

OHTUVAYRE

+! VERONA PHARMA

3MG/2.5ML

N217389 001 Jun 26, 2024

ENTACAPONE

TABLET; ORAL

COMTAN

<u>AB</u>	+	ORION PHARMA	<u>200MG</u>	<u>N020796 001</u>	Oct 19, 1999
-----------	---	--------------	--------------	--------------------	--------------

ENTACAPONE

<u>AB</u>		AJANTA PHARMA LTD	<u>200MG</u>	<u>A205792 001</u>	Aug 31, 2017
<u>AB</u>		ALEMBIC	<u>200MG</u>	<u>A212601 001</u>	Jan 04, 2022

PRESCRIPTION DRUG PRODUCT LIST

ENTACAPONE

TABLET; ORAL

ENTACAPONE

AB	AUROBINDO PHARMA LTD	200MG	A203437 001	Jun 19, 2015
AB	MACLEODS PHARMS LTD	200MG	A207210 001	Jun 05, 2017
AB	SUN PHARM	200MG	A090690 001	Jul 16, 2012
AB	SUNSHINE	200MG	A206669 001	Oct 03, 2018
AB	WOCKHARDT BIO AG	200MG	A078941 001	Aug 16, 2012

ENTECAVIR

SOLUTION; ORAL

BARACLUDE

+	BRISTOL MYERS SQUIBB	0.05MG/ML	N021798 001	Mar 29, 2005
---	----------------------	-----------	-------------	--------------

TABLET; ORAL

BARACLUDE

AB	+	BRISTOL MYERS SQUIBB	0.5MG	N021797 001	Mar 29, 2005
AB	+		1MG	N021797 002	Mar 29, 2005

ENTECAVIR

AB	AMNEAL PHARMS	0.5MG	A206652 001	Nov 12, 2015
AB		1MG	A206652 002	Nov 12, 2015
AB	AUROBINDO PHARMA	0.5MG	A206217 001	Aug 26, 2015
AB		1MG	A206217 002	Aug 26, 2015
AB	BRECKENRIDGE	0.5MG	A208721 001	Mar 15, 2018
AB		1MG	A208721 002	Mar 15, 2018
AB	BRIGHTGENE	0.5MG	A212126 001	Sep 25, 2019
AB		1MG	A212126 002	Sep 25, 2019
AB	CIPLA	0.5MG	A206872 001	Dec 06, 2016
AB		1MG	A206872 002	Dec 06, 2016
AB	CONBA USA	0.5MG	A216857 001	Dec 23, 2024
AB		1MG	A216857 002	Dec 23, 2024
AB	HETERO LABS LTD V	0.5MG	A205740 001	Aug 21, 2015
AB		1MG	A205740 002	Aug 21, 2015
AB	YUNG SHIN PHARM	0.5MG	A208195 001	Nov 10, 2021
AB		1MG	A208195 002	Nov 10, 2021
AB	ZYDUS PHARMS	0.5MG	A206745 001	Jun 23, 2017
AB		1MG	A206745 002	Jun 23, 2017

ENTRECTINIB

CAPSULE; ORAL

ROZLYTREK

+	GENENTECH INC	100MG	N212725 001	Aug 15, 2019
+		200MG	N212725 002	Aug 15, 2019

PELLETS; ORAL

ROZLYTREK

+	GENENTECH INC	50MG/PACKET	N218550 001	Oct 20, 2023
---	---------------	-------------	-------------	--------------

ENZALUTAMIDE

CAPSULE; ORAL

ENZALUTAMIDE

AB	ZYDUS PHARMS	40MG	A209667 001	Sep 26, 2024
-----------	--------------	-------------	--------------------	--------------

XTANDI

AB	+	ASTELLAS	40MG	N203415 001	Aug 31, 2012
-----------	---	----------	-------------	--------------------	--------------

TABLET; ORAL

XTANDI

+	ASTELLAS	40MG	N213674 001	Aug 04, 2020
+		80MG	N213674 002	Aug 04, 2020

EPHEDRINE HYDROCHLORIDE

SOLUTION; INTRAVENOUS

REZIPRES

+	DR REDDYS LABS SA	47MG/10ML (4.7MG/ML)	N213536 004	Dec 07, 2023
---	-------------------	----------------------	-------------	--------------

EPHEDRINE SULFATE

SOLUTION; INTRAVENOUS

AKOVAZ

AP	+	EXELA PHARMA	25MG/5ML (5MG/ML)	N208289 002	Aug 02, 2021
-----------	---	--------------	--------------------------	--------------------	--------------

AP	+		50MG/ML (50MG/ML)	N208289 001	Apr 29, 2016
-----------	---	--	--------------------------	--------------------	--------------

CORPHEDRA

AP	ENDO OPERATIONS	50MG/ML (50MG/ML)	N208943 001	Jan 27, 2017
-----------	-----------------	--------------------------	--------------------	--------------

EPHEDRINE SULFATE

AP	AMNEAL	50MG/ML (50MG/ML)	A212932 001	Oct 23, 2019
-----------	--------	--------------------------	--------------------	--------------

AP	CAPLIN	50MG/ML (50MG/ML)	A219050 001	Jul 18, 2024
-----------	--------	--------------------------	--------------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

EPHEDRINE SULFATE

SOLUTION; INTRAVENOUS

EPHEDRINE SULFATE

<u>AP</u>	DR REDDYS	<u>50MG/ML (50MG/ML)</u>	<u>A212649 001</u>	Oct 03, 2020
<u>AP</u>	EUGIA PHARMA	<u>50MG/ML (50MG/ML)</u>	<u>A214579 001</u>	Jun 14, 2021
<u>AP</u>	FRESENIUS KABI USA	<u>50MG/ML (50MG/ML)</u>	<u>A209646 001</u>	Aug 04, 2020
<u>AP</u>	GLAND PHARMA LTD	<u>50MG/ML (50MG/ML)</u>	<u>A216146 001</u>	Feb 25, 2022
<u>AP</u>	HIKMA	<u>25MG/5ML (5MG/ML)</u>	<u>A217721 001</u>	Oct 11, 2024
<u>AP</u>		<u>50MG/ML (50MG/ML)</u>	<u>A214334 001</u>	Dec 15, 2020
<u>AP</u>	MANKIND PHARMA	<u>50MG/ML (50MG/ML)</u>	<u>A216129 001</u>	Apr 14, 2022
<u>AP</u>	SAGENT PHARMS INC	<u>50MG/ML (50MG/ML)</u>	<u>A214528 001</u>	Mar 09, 2023
<u>AP</u>	SANDOZ	<u>50MG/ML (50MG/ML)</u>	<u>A209784 001</u>	Aug 23, 2017
<u>AP</u>	XIROMED	<u>50MG/ML (50MG/ML)</u>	<u>A215825 001</u>	Apr 21, 2022

EMERPHED

+	!	NEXUS	25MG/5ML (5MG/ML)	N213407 002	Feb 28, 2023
+	!		50MG/10ML (5MG/ML)	N213407 001	Apr 17, 2020

EPHEDRINE SULFATE

+	!	ENDO OPERATIONS	25MG/5ML (5MG/ML)	N213994 002	Apr 22, 2022
+	!		50MG/10ML (5MG/ML)	N213994 001	Oct 16, 2020

EPINASTINE HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

EPINASTINE HYDROCHLORIDE

<u>AT</u>	APOTEX	<u>0.05%</u>	<u>A090919 001</u>	Oct 31, 2011	
<u>AT</u>	!	SOMERSET THERAPS LLC	<u>0.05%</u>	<u>A090951 001</u>	Oct 31, 2011

EPINEPHRINE

INJECTABLE; INTRAMUSCULAR, SUBCUTANEOUS

EPINEPHRINE (AUTOINJECTOR)

<u>AB</u>	TEVA PHARMS USA	<u>0.15MG/DELIVERY</u>	<u>A090589 002</u>	Aug 16, 2018
<u>AB</u>		<u>0.3MG/DELIVERY</u>	<u>A090589 001</u>	Aug 16, 2018

EPIPEN

<u>AB</u>	+	!	MYLAN SPECIALITY LP	<u>0.3MG/DELIVERY</u>	<u>N019430 001</u>	Dec 22, 1987
-----------	---	---	---------------------	-----------------------	--------------------	--------------

EPIPEN JR.

<u>AB</u>	+	!	MYLAN SPECIALITY LP	<u>0.15MG/DELIVERY</u>	<u>N019430 002</u>	Dec 22, 1987
-----------	---	---	---------------------	------------------------	--------------------	--------------

ADRENACLICK

BX	+	!	IMPAX	EQ 0.15MG/DELIVERY	N020800 003	Nov 25, 2009
BX	+	!		EQ 0.3MG/DELIVERY	N020800 004	Nov 25, 2009

SOLUTION; INTRAMUSCULAR, INTRAVENOUS, SUBCUTANEOUS

ADRENALIN

<u>AP</u>	+	!	ENDO OPERATIONS	<u>EQ 30MG BASE/30ML (EQ 1MG BASE/ML)</u>	<u>N204640 001</u>	Dec 18, 2013
-----------	---	---	-----------------	---	--------------------	--------------

EPINEPHRINE

<u>AP</u>	AM REGENT	<u>EQ 30MG BASE/30ML (EQ 1MG BASE/ML)</u>	<u>A217192 001</u>	Sep 06, 2024
<u>AP</u>	INTL MEDICATION SYS	<u>EQ 30MG BASE/30ML (EQ 1MG BASE/ML)</u>	<u>A211880 001</u>	Apr 24, 2020

ADRENALIN

+	!	ENDO OPERATIONS	EQ 1MG BASE/ML (EQ 1MG BASE/ML)	N204200 001	Dec 07, 2012
---	---	-----------------	---------------------------------	-------------	--------------

EPINEPHRINE

+	!	BPI LABS	1MG/ML (1MG/ML)	N205029 005	Mar 04, 2024
+	!		10MG/10ML (1MG/ML)	N205029 002	Feb 04, 2022
+	!		30MG/30ML (1MG/ML)	N205029 004	Feb 14, 2024

SOLUTION; INTRAMUSCULAR, SUBCUTANEOUS

AUVI-Q

BX	+	!	KALEO INC	EQ 0.15MG/DELIVERY	N201739 002	Aug 10, 2012
BX	+			EQ 0.3MG/DELIVERY	N201739 001	Aug 10, 2012
	+			EQ 0.1MG/DELIVERY	N201739 003	Nov 17, 2017

SYMJEPI

+	!	ADAMIS PHARMS CORP	0.3MG/0.3ML (0.3MG/0.3ML)	N207534 001	Jun 15, 2017
---	---	--------------------	---------------------------	-------------	--------------

SOLUTION; INTRAVENOUS

ADRENALIN

+	!	ENDO OPERATIONS	2MG/250ML (8MCG/ML)	N215875 001	Apr 21, 2023
+	!		4MG/250ML (16MCG/ML)	N215875 002	Apr 21, 2023
+	!		5MG/250ML (20MCG/ML)	N215875 003	Apr 21, 2023
+	!		8MG/250ML (32MCG/ML)	N215875 004	Apr 21, 2023
+	!		10MG/250ML (40MCG/ML)	N215875 005	Apr 21, 2023

EPINEPHRINE

+	!	HOSPIRA	1MG/10ML (0.1MG/ML)	N209359 001	Nov 05, 2019
+	!	INTL MEDICATION SYS	1MG/10ML (0.1MG/ML)	N211363 001	Aug 15, 2022

SOLUTION; INTRAVENOUS, INTRAOCULAR, INTRAMUSCULAR, SUBCUTANEOUS

EPINEPHRINE

<u>AP</u>	+	!	BPI LABS	<u>1MG/ML (1MG/ML)</u>	<u>N205029 001</u>	Jul 29, 2014
<u>AP</u>			FRESENIUS KABI USA	<u>1MG/ML (1MG/ML)</u>	<u>A213708 001</u>	Nov 20, 2024

PRESCRIPTION DRUG PRODUCT LIST

EPINEPHRINE

SPRAY;NASAL

NEFFY

+	!	ARS PHARMS	2MG/SPRAY	N214697	001	Aug 09, 2024
		OPERATION				

EPINEPHRINE BITARTRATE; LIDOCAINE HYDROCHLORIDE

INJECTABLE;INJECTION

LIGNOSPAN FORTE

+	!	DEPROCO	EQ 0.02MG BASE/ML;2%	A088389	001	Jan 22, 1985
---	---	---------	----------------------	---------	-----	--------------

LIGNOSPAN STANDARD

+	!	DEPROCO	EQ 0.01MG BASE/ML;2%	A088390	001	Jan 22, 1985
---	---	---------	----------------------	---------	-----	--------------

EPINEPHRINE BITARTRATE; PRILOCAINE HYDROCHLORIDE

INJECTABLE;INJECTION

CITANEST FORTE DENTAL

<u>AP</u>	+	!	DENTSPLY PHARM	<u>0.005MG/ML;4%</u>	<u>N021383</u>	<u>001</u>	
-----------	---	---	----------------	----------------------	----------------	------------	--

PRILOCAINE HYDROCHLORIDE AND EPINEPHRINE BITARTRATE

<u>AP</u>			SEPTODONT INC	<u>0.005MG/ML;4%</u>	<u>A078959</u>	<u>001</u>	Aug 30, 2011
-----------	--	--	---------------	----------------------	----------------	------------	--------------

EPINEPHRINE; LIDOCAINE HYDROCHLORIDE

INJECTABLE;INJECTION

LIDOCAINE HYDROCHLORIDE AND EPINEPHRINE

<u>AP</u>			B BRAUN MEDICAL INC	<u>0.005MG/ML;1.5%</u>	<u>A208475</u>	<u>001</u>	Sep 08, 2021
-----------	--	--	---------------------	------------------------	----------------	------------	--------------

<u>AP</u>			HOSPIRA	<u>0.005MG/ML;0.5%</u>	<u>A089635</u>	<u>001</u>	Jun 21, 1988
-----------	--	--	---------	------------------------	----------------	------------	--------------

<u>AP</u>				<u>0.005MG/ML;1.5%</u>	<u>A088571</u>	<u>001</u>	Sep 13, 1985
-----------	--	--	--	------------------------	----------------	------------	--------------

<u>AP</u>				<u>0.005MG/ML;1.5%</u>	<u>A089645</u>	<u>001</u>	Jun 21, 1988
-----------	--	--	--	------------------------	----------------	------------	--------------

<u>AP</u>				<u>0.005MG/ML;2%</u>	<u>A089651</u>	<u>001</u>	Jun 21, 1988
-----------	--	--	--	----------------------	----------------	------------	--------------

<u>AP</u>				<u>0.01MG/ML;1%</u>	<u>A089644</u>	<u>001</u>	Jun 21, 1988
-----------	--	--	--	---------------------	----------------	------------	--------------

XYLOCAINE W/ EPINEPHRINE

<u>AP</u>	+	!	FRESENIUS KABI USA	<u>0.005MG/ML;0.5%</u>	<u>N006488</u>	<u>012</u>	
-----------	---	---	--------------------	------------------------	----------------	------------	--

<u>AP</u>	+	!		<u>0.005MG/ML;1.5%</u>	<u>N006488</u>	<u>017</u>	
-----------	---	---	--	------------------------	----------------	------------	--

<u>AP</u>	+	!		<u>0.005MG/ML;2%</u>	<u>N006488</u>	<u>019</u>	Nov 13, 1986
-----------	---	---	--	----------------------	----------------	------------	--------------

<u>AP</u>	+	!		<u>0.01MG/ML;1%</u>	<u>N006488</u>	<u>004</u>	
-----------	---	---	--	---------------------	----------------	------------	--

LIDOCAINE HYDROCHLORIDE AND EPINEPHRINE

!			HOSPIRA	0.01MG/ML;2%	A089646	001	Jun 21, 1988
---	--	--	---------	--------------	---------	-----	--------------

XYLOCAINE W/ EPINEPHRINE

+	!		FRESENIUS KABI USA	0.005MG/ML;1%	N006488	018	Nov 13, 1986
---	---	--	--------------------	---------------	---------	-----	--------------

+	!			0.02MG/ML;2%	N006488	005	
---	---	--	--	--------------	---------	-----	--

EPIRUBICIN HYDROCHLORIDE

INJECTABLE;INJECTION

ELLEENCE

<u>AP</u>	+	!	PFIZER INC	<u>200MG/100ML (2MG/ML)</u>	<u>N050778</u>	<u>001</u>	Sep 15, 1999
-----------	---	---	------------	-----------------------------	----------------	------------	--------------

<u>AP</u>	+			<u>50MG/25ML (2MG/ML)</u>	<u>N050778</u>	<u>002</u>	Sep 15, 1999
-----------	---	--	--	---------------------------	----------------	------------	--------------

EPIRUBICIN HYDROCHLORIDE

<u>AP</u>			CIPLA LTD	<u>50MG/25ML (2MG/ML)</u>	<u>A065361</u>	<u>001</u>	Oct 22, 2007
-----------	--	--	-----------	---------------------------	----------------	------------	--------------

<u>AP</u>				<u>200MG/100ML (2MG/ML)</u>	<u>A065361</u>	<u>002</u>	Oct 22, 2007
-----------	--	--	--	-----------------------------	----------------	------------	--------------

<u>AP</u>			EPIC PHARMA LLC	<u>50MG/25ML (2MG/ML)</u>	<u>A090163</u>	<u>001</u>	Jun 24, 2009
-----------	--	--	-----------------	---------------------------	----------------	------------	--------------

<u>AP</u>			HIKMA	<u>50MG/25ML (2MG/ML)</u>	<u>A065289</u>	<u>001</u>	Jun 27, 2007
-----------	--	--	-------	---------------------------	----------------	------------	--------------

<u>AP</u>				<u>200MG/100ML (2MG/ML)</u>	<u>A065289</u>	<u>002</u>	Jun 27, 2007
-----------	--	--	--	-----------------------------	----------------	------------	--------------

<u>AP</u>			HISUN PHARM	<u>50MG/25ML (2MG/ML)</u>	<u>A090075</u>	<u>001</u>	Mar 25, 2010
-----------	--	--	-------------	---------------------------	----------------	------------	--------------

<u>AP</u>			HANGZHOU				
-----------	--	--	----------	--	--	--	--

<u>AP</u>				<u>200MG/100ML (2MG/ML)</u>	<u>A090075</u>	<u>002</u>	Mar 25, 2010
-----------	--	--	--	-----------------------------	----------------	------------	--------------

<u>AP</u>			IMPAX LABS INC	<u>50MG/25ML (2MG/ML)</u>	<u>A065331</u>	<u>001</u>	Aug 09, 2007
-----------	--	--	----------------	---------------------------	----------------	------------	--------------

<u>AP</u>				<u>200MG/100ML (2MG/ML)</u>	<u>A065331</u>	<u>002</u>	Aug 09, 2007
-----------	--	--	--	-----------------------------	----------------	------------	--------------

EPLERENONE

TABLET;ORAL

EPLERENONE

<u>AB</u>			ACCORD HLTHCARE	<u>25MG</u>	<u>A206922</u>	<u>001</u>	Jul 13, 2017
-----------	--	--	-----------------	-------------	----------------	------------	--------------

<u>AB</u>				<u>50MG</u>	<u>A206922</u>	<u>002</u>	Jul 13, 2017
-----------	--	--	--	-------------	----------------	------------	--------------

<u>AB</u>			ADAPTIS	<u>25MG</u>	<u>A212765</u>	<u>001</u>	Aug 10, 2020
-----------	--	--	---------	-------------	----------------	------------	--------------

<u>AB</u>				<u>50MG</u>	<u>A212765</u>	<u>002</u>	Aug 10, 2020
-----------	--	--	--	-------------	----------------	------------	--------------

<u>AB</u>			ANNORA PHARMA	<u>25MG</u>	<u>A213812</u>	<u>001</u>	Jun 02, 2023
-----------	--	--	---------------	-------------	----------------	------------	--------------

<u>AB</u>				<u>50MG</u>	<u>A213812</u>	<u>002</u>	Jun 02, 2023
-----------	--	--	--	-------------	----------------	------------	--------------

<u>AB</u>			BRECKENRIDGE	<u>25MG</u>	<u>A208283</u>	<u>001</u>	Sep 14, 2018
-----------	--	--	--------------	-------------	----------------	------------	--------------

<u>AB</u>				<u>50MG</u>	<u>A208283</u>	<u>002</u>	Sep 14, 2018
-----------	--	--	--	-------------	----------------	------------	--------------

<u>AB</u>			CHARTWELL RX	<u>25MG</u>	<u>A078482</u>	<u>001</u>	Jul 30, 2008
-----------	--	--	--------------	-------------	----------------	------------	--------------

<u>AB</u>				<u>50MG</u>	<u>A078482</u>	<u>002</u>	Jul 30, 2008
-----------	--	--	--	-------------	----------------	------------	--------------

<u>AB</u>			RISING	<u>25MG</u>	<u>A214663</u>	<u>001</u>	Mar 23, 2022
-----------	--	--	--------	-------------	----------------	------------	--------------

<u>AB</u>				<u>50MG</u>	<u>A214663</u>	<u>002</u>	Mar 23, 2022
-----------	--	--	--	-------------	----------------	------------	--------------

<u>AB</u>			SANDOZ	<u>25MG</u>	<u>A078510</u>	<u>001</u>	Aug 01, 2008
-----------	--	--	--------	-------------	----------------	------------	--------------

<u>AB</u>				<u>50MG</u>	<u>A078510</u>	<u>002</u>	Aug 01, 2008
-----------	--	--	--	-------------	----------------	------------	--------------

<u>AB</u>			WESTMINSTER PHARMS	<u>25MG</u>	<u>A207842</u>	<u>001</u>	Oct 25, 2021
-----------	--	--	--------------------	-------------	----------------	------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

EPLERENONE

TABLET; ORAL

EPLERENONE

AB		50MG	A207842 002	Oct 25, 2021
-----------	--	-------------	--------------------	--------------

INSPRA

AB	+	UPJOHN	25MG	N021437 001	Sep 27, 2002
-----------	---	--------	-------------	--------------------	--------------

AB	+	!	50MG	N021437 002	Sep 27, 2002
-----------	---	---	-------------	--------------------	--------------

EPLONTERSEN SODIUM

SOLUTION; SUBCUTANEOUS

WAINUA (AUTOINJECTOR)

+	!	ASTRAZENECA AB	EQ 45MG BASE/0.8ML (EQ 45MG BASE/0.8ML)	N217388 001	Dec 21, 2023
---	---	----------------	---	-------------	--------------

EPOPROSTENOL SODIUM

INJECTABLE; INJECTION

EPOPROSTENOL SODIUM

AP1		MEITHEAL	EQ 0.5MG BASE/VIAL	A078396 001	Apr 23, 2008
------------	--	----------	---------------------------	--------------------	--------------

AP1			EQ 1.5MG BASE/VIAL	A078396 002	Apr 23, 2008
------------	--	--	---------------------------	--------------------	--------------

FLOLAN

AP1	+	!	GLAXOSMITHKLINE LLC	EQ 0.5MG BASE/VIAL	N020444 001	Sep 20, 1995
------------	---	---	---------------------	---------------------------	--------------------	--------------

AP1	+	!		EQ 1.5MG BASE/VIAL	N020444 002	Sep 20, 1995
------------	---	---	--	---------------------------	--------------------	--------------

EPOPROSTENOL SODIUM

AP2		MYLAN	EQ 0.5MG BASE/VIAL	A213913 001	Jun 12, 2024
------------	--	-------	---------------------------	--------------------	--------------

AP2			EQ 1.5MG BASE/VIAL	A213913 002	Jun 12, 2024
------------	--	--	---------------------------	--------------------	--------------

AP2	!	SUN PHARM	EQ 0.5MG BASE/VIAL	A210473 001	Jan 15, 2021
------------	---	-----------	---------------------------	--------------------	--------------

AP2			EQ 1.5MG BASE/VIAL	A210473 002	Jan 15, 2021
------------	--	--	---------------------------	--------------------	--------------

VELETRI

AP2	+	ACTELION	EQ 0.5MG BASE/VIAL	N022260 002	Jun 28, 2012
------------	---	----------	---------------------------	--------------------	--------------

AP2	+	!		EQ 1.5MG BASE/VIAL	N022260 001	Jun 27, 2008
------------	---	---	--	---------------------------	--------------------	--------------

EPTIFIBATIDE

INJECTABLE; INJECTION

EPTIFIBATIDE

AP		BAXTER HLTHCARE CORP	75MG/100ML	A208554 002	Nov 23, 2018
-----------	--	----------------------	-------------------	--------------------	--------------

AP		EUGIA PHARMA	2MG/ML	A206127 001	Dec 08, 2015
-----------	--	--------------	---------------	--------------------	--------------

AP			75MG/100ML	A206127 002	Dec 08, 2015
-----------	--	--	-------------------	--------------------	--------------

AP		HONG KONG	2MG/ML	A213599 001	May 09, 2024
-----------	--	-----------	---------------	--------------------	--------------

AP			75MG/100ML	A213599 002	May 09, 2024
-----------	--	--	-------------------	--------------------	--------------

AP	!	MYLAN LABS LTD	2MG/ML	A203258 001	Jul 20, 2018
-----------	---	----------------	---------------	--------------------	--------------

AP	!		75MG/100ML	A203258 002	Jul 20, 2018
-----------	---	--	-------------------	--------------------	--------------

AP		SAGENT PHARMS INC	2MG/ML	A204693 001	Mar 07, 2018
-----------	--	-------------------	---------------	--------------------	--------------

AP			75MG/100ML	A204693 002	Mar 07, 2018
-----------	--	--	-------------------	--------------------	--------------

AP		SHUANGCHENG	2MG/ML	A213081 001	Sep 07, 2021
-----------	--	-------------	---------------	--------------------	--------------

AP			75MG/100ML	A213081 002	Apr 20, 2022
-----------	--	--	-------------------	--------------------	--------------

AP		SLATE RUN PHARMA	2MG/ML	A209864 001	Jan 25, 2019
-----------	--	------------------	---------------	--------------------	--------------

AP			75MG/100ML	A209864 002	Jan 25, 2019
-----------	--	--	-------------------	--------------------	--------------

AP		TEVA PHARMS USA	2MG/ML	A090854 001	Jun 12, 2015
-----------	--	-----------------	---------------	--------------------	--------------

ERAVACYCLINE DIHYDROCHLORIDE

POWDER; INTRAVENOUS

XERAVA

+	!	TETRAPHASE PHARMS	EQ 50MG BASE/VIAL	N211109 001	Aug 27, 2018
---	---	-------------------	-------------------	-------------	--------------

+	!		EQ 100MG BASE/VIAL	N211109 002	Jun 03, 2020
---	---	--	--------------------	-------------	--------------

ERDAFITINIB

TABLET; ORAL

BALVERSA

+		JANSSEN BIOTECH	3MG	N212018 001	Apr 12, 2019
---	--	-----------------	-----	-------------	--------------

+			4MG	N212018 002	Apr 12, 2019
---	--	--	-----	-------------	--------------

+	!		5MG	N212018 003	Apr 12, 2019
---	---	--	-----	-------------	--------------

ERGOCALCIFEROL

CAPSULE; ORAL

DRISDOL

AA	+	ESJAY PHARMA	50,000 IU	N003444 001	
-----------	---	--------------	------------------	--------------------	--

ERGOCALCIFEROL

AA		CHARTWELL RX	50,000 IU	A040833 001	May 20, 2009
-----------	--	--------------	------------------	--------------------	--------------

AA		PURACAP PHARM LLC	50,000 IU	A204276 001	Dec 07, 2018
-----------	--	-------------------	------------------	--------------------	--------------

AA	!	STRIDES SOFTGELS	50,000 IU	A090455 001	Aug 03, 2010
-----------	---	------------------	------------------	--------------------	--------------

VITAMIN D

AA		BIONPHARMA	50,000 IU	A080704 001	
-----------	--	------------	------------------	--------------------	--

PRESCRIPTION DRUG PRODUCT LIST

ERGOTAMINE TARTRATE

TABLET;SUBLINGUAL

ERGOMAR

! PANGEA

2MG

A087693 001 Feb 24, 1983

ERIBULIN MESYLATE

SOLUTION;INTRAVENOUS

ERIBULIN MESYLATE

<u>AP</u>	BAXTER HLTHCARE CORP	<u>1MG/2ML (0.5MG/ML)</u>	<u>A217250 001</u>	Oct 01, 2024
<u>AP</u>	GLAND PHARMA LTD	<u>1MG/2ML (0.5MG/ML)</u>	<u>A218047 001</u>	Apr 05, 2024
<u>AP</u>	JIANGXI QINGFENG	<u>1MG/2ML (0.5MG/ML)</u>	<u>A218281 001</u>	Jun 28, 2024
<u>AP</u>	LONG GROVE PHARMS	<u>1MG/2ML (0.5MG/ML)</u>	<u>A214850 001</u>	Jul 18, 2024

HALAVEN

<u>AP</u>	+! EISAI INC	<u>1MG/2ML (0.5MG/ML)</u>	<u>N201532 001</u>	Nov 15, 2010
-----------	--------------	---------------------------	--------------------	--------------

ERLOTINIB HYDROCHLORIDE

TABLET;ORAL

ERLOTINIB HYDROCHLORIDE

<u>AB</u>	ALEMbic	<u>EQ 25MG BASE</u>	<u>A214719 001</u>	Jul 08, 2021
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A214719 002</u>	Jul 08, 2021
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A214719 003</u>	Jul 08, 2021
<u>AB</u>	APOTEX	<u>EQ 25MG BASE</u>	<u>A208396 001</u>	Nov 05, 2019
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A208396 002</u>	Nov 05, 2019
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A208396 003</u>	Nov 05, 2019
<u>AB</u>	CHARTWELL RX	<u>EQ 25MG BASE</u>	<u>A203843 001</u>	Sep 13, 2024
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A203843 002</u>	Sep 13, 2024
<u>AB</u>	HETERO LABS LTD V	<u>EQ 25MG BASE</u>	<u>A209267 001</u>	May 24, 2024
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A209267 002</u>	May 24, 2024
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A209267 003</u>	May 24, 2024
<u>AB</u>	MSN	<u>EQ 25MG BASE</u>	<u>A214366 001</u>	May 10, 2021
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A214366 002</u>	May 10, 2021
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A214366 003</u>	May 10, 2021
<u>AB</u>	NATCO PHARMA LTD	<u>EQ 100MG BASE</u>	<u>A208488 002</u>	Nov 05, 2019
<u>AB</u>	RISING	<u>EQ 25MG BASE</u>	<u>A091002 001</u>	Jun 11, 2014
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A091002 002</u>	Jun 11, 2014
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A091002 003</u>	Jun 11, 2014
<u>AB</u>	SHILPA	<u>EQ 25MG BASE</u>	<u>A211960 001</u>	Nov 05, 2019
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A211960 002</u>	Nov 05, 2019
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A211960 003</u>	Nov 05, 2019
<u>AB</u>	SUN PHARM	<u>EQ 25MG BASE</u>	<u>A210300 001</u>	Nov 05, 2019
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A210300 002</u>	Nov 05, 2019
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A210300 003</u>	Nov 05, 2019
<u>AB</u>	TEVA PHARMS USA INC	<u>EQ 25MG BASE</u>	<u>A091059 001</u>	Nov 09, 2020
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A091059 002</u>	Aug 28, 2015
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A091059 003</u>	Aug 28, 2015
<u>AB</u>	ZYDUS PHARMS	<u>EQ 25MG BASE</u>	<u>A213065 001</u>	Apr 16, 2020
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A213065 002</u>	Apr 16, 2020
<u>AB</u>	!	<u>EQ 150MG BASE</u>	<u>A213065 003</u>	Apr 16, 2020

ERTAPENEM SODIUM

INJECTABLE;INTRAMUSCULAR, INTRAVENOUS

ERTAPENEM SODIUM

<u>AP</u>	ACS DOBFAR SPA	<u>EQ 1GM BASE/VIAL</u>	<u>A208790 001</u>	Apr 16, 2018
<u>AP</u>	EUGIA PHARMA	<u>EQ 1GM BASE/VIAL</u>	<u>A209133 001</u>	Jun 25, 2018
<u>AP</u>	GLAND PHARMA LTD	<u>EQ 1GM BASE/VIAL</u>	<u>A212040 001</u>	Mar 26, 2021
<u>AP</u>	QILU ANTIBIOTICS	<u>EQ 1GM BASE/VIAL</u>	<u>A218067 001</u>	Oct 31, 2024
<u>AP</u>	SAVIOR LIFETEC CORP	<u>EQ 1GM BASE/VIAL</u>	<u>A207647 001</u>	Mar 19, 2019

INVANZ

<u>AP</u>	+! MSD SUB MERCK	<u>EQ 1GM BASE/VIAL</u>	<u>N021337 001</u>	Nov 21, 2001
-----------	------------------	-------------------------	--------------------	--------------

ERTUGLIFLOZIN

TABLET;ORAL

ERTUGLIFLOZIN

<u>AB</u>	HETERO LABS LTD V	<u>5MG</u>	<u>A217071 001</u>	Sep 17, 2024
<u>AB</u>		<u>15MG</u>	<u>A217071 002</u>	Sep 17, 2024

STEGLATRO

<u>AB</u>	+ MSD SUB MERCK	<u>5MG</u>	<u>N209803 001</u>	Dec 19, 2017
<u>AB</u>	+!	<u>15MG</u>	<u>N209803 002</u>	Dec 19, 2017

PRESCRIPTION DRUG PRODUCT LIST

ERTUGLIFLOZIN; METFORMIN HYDROCHLORIDE

TABLET;ORAL

SEGLUROMET

+	MSD SUB MERCK	2.5MG;500MG	N209806 001	Dec 19, 2017
+		2.5MG;1GM	N209806 002	Dec 19, 2017
+		7.5MG;500MG	N209806 003	Dec 19, 2017
+	!	7.5MG;1GM	N209806 004	Dec 19, 2017

ERTUGLIFLOZIN; SITAGLIPTIN PHOSPHATE

TABLET;ORAL

STEGLUJAN

+	MSD SUB MERCK	5MG;EQ 100MG BASE	N209805 001	Dec 19, 2017
+	!	15MG;EQ 100MG BASE	N209805 002	Dec 19, 2017

ERYTHROMYCIN

CAPSULE, DELAYED REL PELLETS;ORAL

ERYC

AB	+ !	DR REDDYS LABS SA	250MG	N050536 001	
AB		CARNEGIE	250MG	A062746 001	Dec 22, 1986

GEL;TOPICAL

ERYGEL

AT	+ !	MYLAN	2%	N050617 001	Oct 21, 1987
AT		FOUGERA PHARMS	2%	A064184 001	Sep 30, 1997
AT		PADAGIS US	2%	A063211 001	Jan 29, 1993

OINTMENT;OPHTHALMIC

ERYTHROMYCIN

AT	!	BAUSCH AND LOMB	0.5%	A064067 001	Jul 29, 1994
AT		SENTISS	0.5%	A064030 001	Jul 18, 1996

SOLUTION;TOPICAL

ERYTHRA-DERM

AT		MICRO LABS	2%	A062687 001	Feb 05, 1988
AT	!	PADAGIS US	2%	A063038 001	Jan 11, 1991

SWAB;TOPICAL

ERYTHROMYCIN

AT		EPIC PHARMA LLC	2%	A090215 001	May 12, 2010
AT	!	PADAGIS US	2%	A064126 001	Jul 03, 1996

TABLET;ORAL

ERYTHROMYCIN

AB		ALEMBIC	250MG	A215661 001	Aug 24, 2023
AB			500MG	A215661 002	Aug 24, 2023
AB		ALKEM LABS LTD	250MG	A216066 001	Jul 13, 2022
AB			500MG	A216066 002	Jul 13, 2022
AB		CADILA PHARMS LTD	250MG	A213628 001	Jun 28, 2021
AB	!		500MG	A213628 002	Jun 28, 2021
AB		CARNEGIE	250MG	A061621 001	
AB			500MG	A061621 002	
AB		TEVA PHARMS USA INC	250MG	A214549 001	Feb 11, 2021
AB			500MG	A214549 002	Feb 11, 2021
AB		TORRENT	250MG	A212015 001	Jul 06, 2020
AB			500MG	A212015 002	Jul 06, 2020
AB		ZYDUS LIFESCIENCES	250MG	A212693 001	Mar 07, 2023
AB			500MG	A212693 002	Mar 07, 2023

TABLET, DELAYED RELEASE;ORAL

ERY-TAB

AB		CARNEGIE	250MG	A062298 001	
AB			333MG	A062298 003	Mar 29, 1982
AB	!		500MG	A062298 002	

ERYTHROMYCIN

AB		AMNEAL PHARMS CO	250MG	A210954 001	Jul 02, 2019
AB			333MG	A210954 002	Jul 02, 2019
AB			500MG	A210954 003	Jul 02, 2019
AB		TORRENT	250MG	A211975 001	Jul 26, 2021
AB			333MG	A211975 003	Aug 13, 2024
AB			500MG	A211975 002	Jul 26, 2021

PRESCRIPTION DRUG PRODUCT LIST

ERYTHROMYCIN ETHYLSUCCINATE

GRANULE; ORAL

E.E.S.

<u>AB</u>	+	CARNEGIE	<u>EQ 200MG BASE/5ML</u>	<u>N050207 001</u>	
-----------	---	----------	--------------------------	--------------------	--

ERYPED

<u>AB</u>	+	CARNEGIE	<u>EQ 200MG BASE/5ML</u>	<u>N050207 003</u>	Mar 30, 1987
-----------	---	----------	--------------------------	--------------------	--------------

<u>AB</u>	+	!	<u>EQ 400MG BASE/5ML</u>	<u>N050207 002</u>	
-----------	---	---	--------------------------	--------------------	--

ERYTHROMYCIN ETHYLSUCCINATE

<u>AB</u>		AMNEAL PHARMS	<u>EQ 200MG BASE/5ML</u>	<u>A211204 001</u>	Nov 01, 2019
-----------	--	---------------	--------------------------	--------------------	--------------

<u>AB</u>			<u>EQ 400MG BASE/5ML</u>	<u>A211204 002</u>	Nov 01, 2019
-----------	--	--	--------------------------	--------------------	--------------

<u>AB</u>		ANI PHARMS	<u>EQ 200MG BASE/5ML</u>	<u>A062055 001</u>	
-----------	--	------------	--------------------------	--------------------	--

<u>AB</u>			<u>EQ 200MG BASE/5ML</u>	<u>A062055 003</u>	Nov 02, 2018
-----------	--	--	--------------------------	--------------------	--------------

<u>AB</u>			<u>EQ 400MG BASE/5ML</u>	<u>A062055 002</u>	Nov 02, 2018
-----------	--	--	--------------------------	--------------------	--------------

<u>AB</u>		CADILA PHARMS LTD	<u>EQ 200MG BASE/5ML</u>	<u>A216212 001</u>	Nov 21, 2022
-----------	--	-------------------	--------------------------	--------------------	--------------

<u>AB</u>			<u>EQ 400MG BASE/5ML</u>	<u>A216212 002</u>	Nov 21, 2022
-----------	--	--	--------------------------	--------------------	--------------

TABLET; ORAL

E.E.S. 400

<u>BX</u>	!	CARNEGIE	<u>EQ 400MG BASE</u>	<u>A061905 002</u>	Aug 12, 1982
-----------	---	----------	----------------------	--------------------	--------------

ERYTHROMYCIN ETHYLSUCCINATE

<u>BX</u>	!	CARNEGIE	<u>EQ 400MG BASE</u>	<u>A061904 001</u>	
-----------	---	----------	----------------------	--------------------	--

ERYTHROMYCIN LACTOBIONATE

INJECTABLE; INJECTION

ERYTHROCIN

<u>AP</u>		HOSPIRA	<u>EQ 500MG BASE/VIAL</u>	<u>A062638 001</u>	Oct 31, 1986
-----------	--	---------	---------------------------	--------------------	--------------

<u>AP</u>	+	!	<u>EQ 500MG BASE/VIAL</u>	<u>N050609 001</u>	Sep 24, 1986
-----------	---	---	---------------------------	--------------------	--------------

ERYTHROMYCIN LACTOBIONATE

<u>AP</u>		GLAND PHARMA LTD	<u>EQ 500MG BASE/VIAL</u>	<u>A216761 001</u>	Nov 19, 2024
-----------	--	------------------	---------------------------	--------------------	--------------

<u>AP</u>		NEXUS	<u>EQ 500MG BASE/VIAL</u>	<u>A215290 001</u>	Feb 14, 2022
-----------	--	-------	---------------------------	--------------------	--------------

ERYTHROMYCIN STEARATE

TABLET; ORAL

ERYTHROCIN STEARATE

!

AZURITY

<u>EQ 250MG BASE</u>	<u>A060359 001</u>	
----------------------	--------------------	--

ESCITALOPRAM OXALATE

SOLUTION; ORAL

ESCITALOPRAM OXALATE

<u>AA</u>		AMNEAL PHARMS	<u>EQ 5MG BASE/5ML</u>	<u>A202227 001</u>	Mar 14, 2012
-----------	--	---------------	------------------------	--------------------	--------------

<u>AA</u>		AUROBINDO PHARMA LTD	<u>EQ 5MG BASE/5ML</u>	<u>A079062 001</u>	Apr 02, 2012
-----------	--	----------------------	------------------------	--------------------	--------------

<u>AA</u>	!	CHARTWELL MOLECULAR	<u>EQ 5MG BASE/5ML</u>	<u>A090477 001</u>	Jun 12, 2013
-----------	---	---------------------	------------------------	--------------------	--------------

<u>AA</u>	!	HETERO LABS LTD III	<u>EQ 5MG BASE/5ML</u>	<u>A202221 001</u>	Jun 12, 2012
-----------	---	---------------------	------------------------	--------------------	--------------

<u>AA</u>		MACLEODS PHARMS LTD	<u>EQ 5MG BASE/5ML</u>	<u>A202754 001</u>	Mar 31, 2016
-----------	--	---------------------	------------------------	--------------------	--------------

<u>AA</u>		TARO	<u>EQ 5MG BASE/5ML</u>	<u>A079121 001</u>	May 03, 2012
-----------	--	------	------------------------	--------------------	--------------

TABLET; ORAL

ESCITALOPRAM OXALATE

<u>AB</u>		ACCORD HLTHCARE	<u>EQ 5MG BASE</u>	<u>A202389 001</u>	Sep 11, 2012
-----------	--	-----------------	--------------------	--------------------	--------------

<u>AB</u>			<u>EQ 10MG BASE</u>	<u>A202389 002</u>	Sep 11, 2012
-----------	--	--	---------------------	--------------------	--------------

<u>AB</u>			<u>EQ 20MG BASE</u>	<u>A202389 003</u>	Sep 11, 2012
-----------	--	--	---------------------	--------------------	--------------

<u>AB</u>		AMNEAL PHARMS	<u>EQ 5MG BASE</u>	<u>A205619 001</u>	May 17, 2017
-----------	--	---------------	--------------------	--------------------	--------------

<u>AB</u>			<u>EQ 10MG BASE</u>	<u>A205619 002</u>	May 17, 2017
-----------	--	--	---------------------	--------------------	--------------

<u>AB</u>			<u>EQ 20MG BASE</u>	<u>A205619 003</u>	May 17, 2017
-----------	--	--	---------------------	--------------------	--------------

<u>AB</u>		AUROBINDO PHARMA LTD	<u>EQ 5MG BASE</u>	<u>A090432 001</u>	Sep 11, 2012
-----------	--	----------------------	--------------------	--------------------	--------------

<u>AB</u>			<u>EQ 10MG BASE</u>	<u>A090432 002</u>	Sep 11, 2012
-----------	--	--	---------------------	--------------------	--------------

<u>AB</u>			<u>EQ 20MG BASE</u>	<u>A090432 003</u>	Sep 11, 2012
-----------	--	--	---------------------	--------------------	--------------

<u>AB</u>		CHARTWELL RX	<u>EQ 5MG BASE</u>	<u>A078169 001</u>	Sep 11, 2012
-----------	--	--------------	--------------------	--------------------	--------------

<u>AB</u>			<u>EQ 10MG BASE</u>	<u>A078169 002</u>	Sep 11, 2012
-----------	--	--	---------------------	--------------------	--------------

<u>AB</u>			<u>EQ 20MG BASE</u>	<u>A078169 003</u>	Sep 11, 2012
-----------	--	--	---------------------	--------------------	--------------

<u>AB</u>		GRAVITI PHARMS	<u>EQ 5MG BASE</u>	<u>A078777 001</u>	Sep 11, 2012
-----------	--	----------------	--------------------	--------------------	--------------

<u>AB</u>			<u>EQ 10MG BASE</u>	<u>A078777 002</u>	Sep 11, 2012
-----------	--	--	---------------------	--------------------	--------------

<u>AB</u>			<u>EQ 20MG BASE</u>	<u>A078777 003</u>	Sep 11, 2012
-----------	--	--	---------------------	--------------------	--------------

<u>AB</u>		INVAGEN PHARMS	<u>EQ 5MG BASE</u>	<u>A078604 001</u>	Sep 11, 2012
-----------	--	----------------	--------------------	--------------------	--------------

<u>AB</u>			<u>EQ 10MG BASE</u>	<u>A078604 002</u>	Sep 11, 2012
-----------	--	--	---------------------	--------------------	--------------

<u>AB</u>			<u>EQ 20MG BASE</u>	<u>A078604 003</u>	Sep 11, 2012
-----------	--	--	---------------------	--------------------	--------------

<u>AB</u>		JUBILANT CADISTA	<u>EQ 5MG BASE</u>	<u>A202280 001</u>	Sep 12, 2012
-----------	--	------------------	--------------------	--------------------	--------------

<u>AB</u>			<u>EQ 10MG BASE</u>	<u>A202280 002</u>	Sep 12, 2012
-----------	--	--	---------------------	--------------------	--------------

<u>AB</u>			<u>EQ 20MG BASE</u>	<u>A202280 003</u>	Sep 12, 2012
-----------	--	--	---------------------	--------------------	--------------

<u>AB</u>		MACLEODS PHARMS LTD	<u>EQ 5MG BASE</u>	<u>A202210 001</u>	Sep 11, 2012
-----------	--	---------------------	--------------------	--------------------	--------------

<u>AB</u>			<u>EQ 10MG BASE</u>	<u>A202210 002</u>	Sep 11, 2012
-----------	--	--	---------------------	--------------------	--------------

<u>AB</u>			<u>EQ 20MG BASE</u>	<u>A202210 003</u>	Sep 11, 2012
-----------	--	--	---------------------	--------------------	--------------

<u>AB</u>		PRINSTON INC	<u>EQ 5MG BASE</u>	<u>A078032 001</u>	Aug 28, 2015
-----------	--	--------------	--------------------	--------------------	--------------

<u>AB</u>			<u>EQ 10MG BASE</u>	<u>A078032 002</u>	Aug 28, 2015
-----------	--	--	---------------------	--------------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

ESCITALOPRAM OXALATE

TABLET; ORAL

ESCITALOPRAM OXALATE

<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A078032 003</u>	Aug 28, 2015
<u>AB</u>	TORRENT PHARMS LTD	<u>EQ 5MG BASE</u>	<u>A090939 001</u>	Sep 11, 2012
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A090939 002</u>	Sep 11, 2012
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A090939 003</u>	Sep 11, 2012
<u>AB</u>	ZYDUS LIFESCIENCES	<u>EQ 5MG BASE</u>	<u>A077734 001</u>	Sep 11, 2012
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A077734 002</u>	Sep 11, 2012
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A077734 003</u>	Sep 11, 2012

LEXAPRO

<u>AB</u>	+	ABBVIE	<u>EQ 5MG BASE</u>	<u>N021323 001</u>	Aug 14, 2002
<u>AB</u>	+		<u>EQ 10MG BASE</u>	<u>N021323 002</u>	Aug 14, 2002
<u>AB</u>	+		<u>EQ 20MG BASE</u>	<u>N021323 003</u>	Aug 14, 2002

ESKETAMINE HYDROCHLORIDE

SPRAY; NASAL

SPRAVATO

+	JANSSEN PHARMS	EQ 28MG BASE	N211243 001	Mar 05, 2019
---	----------------	--------------	-------------	--------------

ESLICARBAZEPINE ACETATE

TABLET; ORAL

APTIOM

<u>AB</u>	+	SUMITOMO PHARMA AM	<u>200MG</u>	<u>N022416 001</u>	Nov 08, 2013
<u>AB</u>	+		<u>400MG</u>	<u>N022416 002</u>	Nov 08, 2013
<u>AB</u>	+		<u>600MG</u>	<u>N022416 003</u>	Nov 08, 2013
<u>AB</u>	+		<u>800MG</u>	<u>N022416 004</u>	Nov 08, 2013

ESLICARBAZEPINE ACETATE

<u>AB</u>		ALKEM LABS LTD	<u>200MG</u>	<u>A211199 001</u>	Oct 06, 2023
<u>AB</u>			<u>400MG</u>	<u>A211199 002</u>	Oct 06, 2023
<u>AB</u>			<u>600MG</u>	<u>A211199 003</u>	Oct 06, 2023
<u>AB</u>			<u>800MG</u>	<u>A211199 004</u>	Oct 06, 2023
<u>AB</u>		DR REDDYS	<u>200MG</u>	<u>A211238 001</u>	Jun 29, 2021
<u>AB</u>			<u>400MG</u>	<u>A211238 002</u>	Jun 29, 2021
<u>AB</u>			<u>600MG</u>	<u>A211238 003</u>	Jun 29, 2021
<u>AB</u>			<u>800MG</u>	<u>A211238 004</u>	Jun 29, 2021
<u>AB</u>		HETERO LABS LTD V	<u>200MG</u>	<u>A211186 001</u>	Aug 03, 2023
<u>AB</u>			<u>400MG</u>	<u>A211186 002</u>	Aug 03, 2023
<u>AB</u>			<u>600MG</u>	<u>A211186 003</u>	Aug 03, 2023
<u>AB</u>			<u>800MG</u>	<u>A211186 004</u>	Aug 03, 2023

ESMOLOL HYDROCHLORIDE

INJECTABLE; INJECTION

BREVIBLOC

<u>AP</u>	+	BAXTER HLTHCARE	<u>10MG/ML</u>	<u>N019386 006</u>	Feb 25, 2003
-----------	---	-----------------	----------------	--------------------	--------------

BREVIBLOC DOUBLE STRENGTH IN PLASTIC CONTAINER

<u>AP</u>	+	BAXTER HLTHCARE	<u>2GM/100ML</u>	<u>N019386 005</u>	Jan 27, 2003
-----------	---	-----------------	------------------	--------------------	--------------

BREVIBLOC IN PLASTIC CONTAINER

<u>AP</u>	+	BAXTER HLTHCARE	<u>1GM/100ML</u>	<u>N019386 004</u>	Feb 16, 2001
-----------	---	-----------------	------------------	--------------------	--------------

ESMOLOL HYDROCHLORIDE

<u>AP</u>		AMNEAL	<u>1GM/100ML</u>	<u>A216603 001</u>	Dec 13, 2022
<u>AP</u>			<u>2GM/100ML</u>	<u>A216603 002</u>	Dec 13, 2022
<u>AP</u>		EUGIA PHARMA	<u>10MG/ML</u>	<u>A205520 001</u>	Jul 23, 2015
<u>AP</u>			<u>1GM/100ML</u>	<u>A216244 001</u>	Mar 21, 2022
<u>AP</u>			<u>2GM/100ML</u>	<u>A216244 002</u>	Mar 21, 2022
<u>AP</u>		GLAND PHARMA LTD	<u>10MG/ML</u>	<u>A208538 001</u>	Aug 14, 2019
<u>AP</u>		HIKMA	<u>10MG/ML</u>	<u>A076323 001</u>	Aug 10, 2004
<u>AP</u>		HQ SPCLT PHARMA	<u>1GM/100ML</u>	<u>A214172 001</u>	Dec 02, 2022
<u>AP</u>			<u>2GM/100ML</u>	<u>A214172 002</u>	Dec 02, 2022
<u>AP</u>		MYLAN INSTITUTIONAL	<u>10MG/ML</u>	<u>A076474 001</u>	May 02, 2005
<u>AP</u>		MYLAN LABS LTD	<u>1GM/100ML</u>	<u>A206608 001</u>	Jun 08, 2018
<u>AP</u>			<u>2GM/100ML</u>	<u>A206608 002</u>	Jun 08, 2018
<u>AP</u>		SAGENT PHARMS INC	<u>1GM/100ML</u>	<u>A207107 001</u>	Jun 08, 2018
<u>AP</u>			<u>2GM/100ML</u>	<u>A207107 002</u>	Jun 08, 2018

SOLUTION; INTRAVENOUS

ESMOLOL HYDROCHLORIDE DOUBLE STRENGTH IN PLASTIC CONTAINER

+	HQ SPCLT PHARMA	2GM/100ML (20MG/ML)	N205703 002	Apr 07, 2016
---	-----------------	---------------------	-------------	--------------

ESMOLOL HYDROCHLORIDE IN PLASTIC CONTAINER

+	HQ SPCLT PHARMA	2.5GM/250ML (10MG/ML)	N205703 001	Apr 07, 2016
---	-----------------	-----------------------	-------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

ESOMEPRAZOLE MAGNESIUM

CAPSULE, DELAYED REL PELLETS;ORAL

ESOMEPRAZOLE MAGNESIUM

<u>AB</u>	ALKEM LABS LTD	<u>EQ 20MG BASE</u>	<u>A208333 001</u>	Oct 20, 2017
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A208333 002</u>	Oct 20, 2017
<u>AB</u>	AUROBINDO PHARMA LTD	<u>EQ 20MG BASE</u>	<u>A205606 001</u>	Apr 21, 2016
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A205606 002</u>	Apr 21, 2016
<u>AB</u>	CISEN	<u>EQ 20MG BASE</u>	<u>A213158 001</u>	Sep 22, 2020
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A213158 002</u>	Sep 22, 2020
<u>AB</u>	CSPC OUYI	<u>EQ 20MG BASE</u>	<u>A212949 001</u>	Oct 02, 2020
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A212949 002</u>	Oct 02, 2020
<u>AB</u>	DR REDDYS	<u>EQ 20MG BASE</u>	<u>A078279 001</u>	Sep 25, 2015
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A078279 002</u>	Sep 25, 2015
<u>AB</u>	ETHYPHARM	<u>EQ 20MG BASE</u>	<u>A090841 001</u>	Mar 31, 2021
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A090841 002</u>	Mar 31, 2021
<u>AB</u>	GRANULES	<u>EQ 20MG BASE</u>	<u>A217427 001</u>	Oct 18, 2023
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A217427 002</u>	Oct 18, 2023
<u>AB</u>	GRAVITI PHARMS	<u>EQ 20MG BASE</u>	<u>A213486 001</u>	Mar 19, 2021
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A213486 002</u>	Mar 19, 2021
<u>AB</u>	GUANGZHOU NOVAKEN	<u>EQ 20MG BASE</u>	<u>A213859 001</u>	Nov 18, 2020
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A213859 002</u>	Nov 18, 2020
<u>AB</u>	HETERO LABS LTD III	<u>EQ 20MG BASE</u>	<u>A211977 001</u>	Jun 02, 2020
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A211977 002</u>	Jun 02, 2020
<u>AB</u>	INDCHEMIE HEALTH	<u>EQ 20MG BASE</u>	<u>A210559 001</u>	Feb 26, 2021
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A210559 002</u>	Feb 26, 2021
<u>AB</u>	IVAX SUB TEVA PHARMS	<u>EQ 20MG BASE</u>	<u>A078003 001</u>	Jan 26, 2015
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A078003 002</u>	Jan 26, 2015
<u>AB</u>	LANNETT CO INC	<u>EQ 20MG BASE</u>	<u>A205563 001</u>	Sep 01, 2017
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A205563 002</u>	Sep 01, 2017
<u>AB</u>	MACLEODS PHARMS LTD	<u>EQ 20MG BASE</u>	<u>A208511 001</u>	Oct 23, 2024
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A208511 002</u>	Oct 23, 2024
<u>AB</u>	MYLAN	<u>EQ 20MG BASE</u>	<u>A078936 001</u>	Aug 02, 2015
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A078936 002</u>	Aug 03, 2015
<u>AB</u>	PRINSTON INC	<u>EQ 20MG BASE</u>	<u>A214920 001</u>	Mar 28, 2023
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A214920 002</u>	Mar 28, 2023
<u>AB</u>	ZHEJIANG YONGTAI	<u>EQ 20MG BASE</u>	<u>A217022 001</u>	Dec 27, 2023
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A217022 002</u>	Dec 27, 2023
<u>AB</u>	ZYDUS PHARMS	<u>EQ 20MG BASE</u>	<u>A206296 001</u>	May 22, 2019
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A206296 002</u>	May 22, 2019

NEXIUM

<u>AB</u>	+ ASTRAZENECA	<u>EQ 20MG BASE</u>	<u>N021153 001</u>	Feb 20, 2001
<u>AB</u>	+!	<u>EQ 40MG BASE</u>	<u>N021153 002</u>	Feb 20, 2001

FOR SUSPENSION, DELAYED RELEASE;ORAL

ESOMEPRAZOLE MAGNESIUM

<u>AB</u>	CIPLA	<u>EQ 2.5MG BASE/PACKET</u>	<u>A217714 001</u>	Jan 07, 2025
<u>AB</u>		<u>EQ 5MG BASE/PACKET</u>	<u>A217714 002</u>	Jan 07, 2025
<u>AB</u>		<u>EQ 10MG BASE/PACKET</u>	<u>A211752 001</u>	Mar 23, 2020
<u>AB</u>		<u>EQ 20MG BASE/PACKET</u>	<u>A211751 001</u>	Mar 23, 2020
<u>AB</u>		<u>EQ 40MG BASE/PACKET</u>	<u>A211751 002</u>	Mar 23, 2020
<u>AB</u>	ZYDUS PHARMS	<u>EQ 20MG BASE/PACKET</u>	<u>A206055 001</u>	Jun 07, 2023
<u>AB</u>		<u>EQ 40MG BASE/PACKET</u>	<u>A206055 002</u>	Jun 07, 2023

NEXIUM

<u>AB</u>	+ ASTRAZENECA	<u>EQ 2.5MG BASE/PACKET</u>	<u>N021957 003</u>	Dec 15, 2011
<u>AB</u>	+	<u>EQ 5MG BASE/PACKET</u>	<u>N021957 004</u>	Dec 15, 2011
<u>AB</u>	+	<u>EQ 10MG BASE/PACKET</u>	<u>N022101 001</u>	Feb 27, 2008
<u>AB</u>	+	<u>EQ 20MG BASE/PACKET</u>	<u>N021957 001</u>	Oct 20, 2006
<u>AB</u>	+!	<u>EQ 40MG BASE/PACKET</u>	<u>N021957 002</u>	Oct 20, 2006

ESOMEPRAZOLE MAGNESIUM; NAPROXEN

TABLET, DELAYED RELEASE;ORAL

NAPROXEN AND ESOMEPRAZOLE MAGNESIUM

<u>AB</u>	AJANTA PHARMA LTD	<u>EQ 20MG BASE;375MG</u>	<u>A213699 001</u>	Oct 06, 2022
<u>AB</u>		<u>EQ 20MG BASE;500MG</u>	<u>A213699 002</u>	Oct 06, 2022
<u>AB</u>	DR REDDYS	<u>EQ 20MG BASE;375MG</u>	<u>A204206 001</u>	Feb 18, 2020
<u>AB</u>	!	<u>EQ 20MG BASE;500MG</u>	<u>A204206 002</u>	Feb 18, 2020
<u>AB</u>	SCIEGEN PHARMS INC	<u>EQ 20MG BASE;375MG</u>	<u>A217738 001</u>	Oct 11, 2023
<u>AB</u>		<u>EQ 20MG BASE;500MG</u>	<u>A217738 002</u>	Oct 11, 2023

PRESCRIPTION DRUG PRODUCT LIST

ESOMEPRAZOLE SODIUM

INJECTABLE; INTRAVENOUS

ESOMEPRAZOLE SODIUM

<u>AP</u>	DEVA HOLDING AS	<u>EQ 40MG BASE/VIAL</u>	<u>A207181 001</u>	Mar 06, 2017
<u>AP</u>	EUGIA PHARMA	<u>EQ 40MG BASE/VIAL</u>	<u>A204657 002</u>	Aug 10, 2016
<u>AP</u>	GLAND PHARMA LTD	<u>EQ 40MG BASE/VIAL</u>	<u>A203349 002</u>	Apr 01, 2020
<u>AP</u>	! SLATE RUN PHARMA	<u>EQ 40MG BASE/VIAL</u>	<u>A215732 001</u>	Feb 10, 2022
	! GLAND PHARMA LTD	EQ 20MG BASE/VIAL	A203349 001	Apr 01, 2020

ESTAZOLAM

TABLET; ORAL

ESTAZOLAM

<u>AB</u>	DR REDDYS LABS SA	<u>1MG</u>	<u>A074921 001</u>	Jul 10, 1997
<u>AB</u>	!	<u>2MG</u>	<u>A074921 002</u>	Jul 10, 1997
<u>AB</u>	NOVITIUM PHARMA	<u>1MG</u>	<u>A074826 001</u>	Jul 03, 1997
<u>AB</u>		<u>2MG</u>	<u>A074826 002</u>	Jul 03, 1997
<u>AB</u>	WATSON LABS	<u>1MG</u>	<u>A074818 001</u>	Aug 19, 1997
<u>AB</u>		<u>2MG</u>	<u>A074818 002</u>	Aug 19, 1997

ESTRADIOL

CREAM; VAGINAL

ESTRACE

<u>AB</u>	+! ALLERGAN	<u>0.01%</u>	<u>A086069 001</u>	Jan 31, 1984
-----------	-------------	--------------	--------------------	--------------

ESTRADIOL

<u>AB</u>	ALVOGEN	<u>0.01%</u>	<u>A209767 001</u>	Mar 05, 2018
<u>AB</u>	MYLAN	<u>0.01%</u>	<u>A208788 001</u>	Dec 29, 2017
<u>AB</u>	PADAGIS ISRAEL	<u>0.01%</u>	<u>A210194 001</u>	Jan 22, 2018
<u>AB</u>	PRASCO	<u>0.01%</u>	<u>A212313 001</u>	Jul 15, 2021
<u>AB</u>	TEVA PHARMS USA	<u>0.01%</u>	<u>A210488 001</u>	Mar 30, 2018

FILM, EXTENDED RELEASE; TRANSDERMAL

CLIMARA

<u>AB2</u>	+ BAYER HLTHCARE	<u>0.025MG/24HR</u>	<u>N020375 004</u>	Mar 05, 1999
<u>AB2</u>	+	<u>0.0375MG/24HR</u>	<u>N020375 005</u>	May 27, 2003
<u>AB2</u>	+	<u>0.05MG/24HR</u>	<u>N020375 001</u>	Dec 22, 1994
<u>AB2</u>	+	<u>0.06MG/24HR</u>	<u>N020375 006</u>	May 27, 2003
<u>AB2</u>	+	<u>0.075MG/24HR</u>	<u>N020375 003</u>	Mar 23, 1998
<u>AB2</u>	+!	<u>0.1MG/24HR</u>	<u>N020375 002</u>	Dec 22, 1994

ESTRADIOL

<u>AB2</u>	MYLAN TECHNOLOGIES	<u>0.025MG/24HR</u>	<u>A075182 003</u>	Jan 26, 2005
<u>AB2</u>		<u>0.0375MG/24HR</u>	<u>A075182 004</u>	Jul 20, 2006
<u>AB2</u>		<u>0.05MG/24HR</u>	<u>A075182 006</u>	Feb 24, 2000
<u>AB2</u>		<u>0.06MG/24HR</u>	<u>A075182 005</u>	Jul 20, 2006
<u>AB2</u>		<u>0.075MG/24HR</u>	<u>A075182 002</u>	Jan 26, 2005
<u>AB2</u>		<u>0.1MG/24HR</u>	<u>A075182 001</u>	Feb 24, 2000
<u>AB2</u>	ZYDUS PHARMS	<u>0.025MG/24HR</u>	<u>A202985 001</u>	Mar 29, 2023
<u>AB2</u>		<u>0.0375MG/24HR</u>	<u>A202985 002</u>	Mar 29, 2023
<u>AB2</u>		<u>0.05MG/24HR</u>	<u>A202985 003</u>	Mar 29, 2023
<u>AB2</u>		<u>0.06MG/24HR</u>	<u>A202985 004</u>	Mar 29, 2023
<u>AB2</u>		<u>0.075MG/24HR</u>	<u>A202985 005</u>	Mar 29, 2023
<u>AB2</u>		<u>0.1MG/24HR</u>	<u>A202985 006</u>	Mar 29, 2023
<u>AB3</u>	AMNEAL	<u>0.025MG/24HR</u>	<u>A211396 001</u>	Sep 28, 2020
<u>AB3</u>		<u>0.0375MG/24HR</u>	<u>A211396 002</u>	Sep 28, 2020
<u>AB3</u>		<u>0.05MG/24HR</u>	<u>A211396 003</u>	Sep 28, 2020
<u>AB3</u>		<u>0.075MG/24HR</u>	<u>A211396 004</u>	Sep 28, 2020
<u>AB3</u>		<u>0.1MG/24HR</u>	<u>A211396 005</u>	Sep 28, 2020
<u>AB3</u>	MYLAN TECHNOLOGIES	<u>0.025MG/24HR</u>	<u>A206685 001</u>	Aug 15, 2018
<u>AB3</u>		<u>0.0375MG/24HR</u>	<u>A206685 002</u>	Aug 15, 2018
<u>AB3</u>		<u>0.05MG/24HR</u>	<u>A206685 003</u>	Aug 15, 2018
<u>AB3</u>		<u>0.075MG/24HR</u>	<u>A206685 004</u>	Aug 15, 2018
<u>AB3</u>		<u>0.1MG/24HR</u>	<u>A206685 005</u>	Aug 15, 2018

MINIVELLE

<u>AB3</u>	+ NOVEN	<u>0.025MG/24HR</u>	<u>N203752 005</u>	Sep 23, 2014
<u>AB3</u>	+	<u>0.0375MG/24HR</u>	<u>N203752 001</u>	Oct 29, 2012
<u>AB3</u>	+	<u>0.05MG/24HR</u>	<u>N203752 003</u>	Oct 29, 2012
<u>AB3</u>	+	<u>0.075MG/24HR</u>	<u>N203752 002</u>	Oct 29, 2012
<u>AB3</u>	+!	<u>0.1MG/24HR</u>	<u>N203752 004</u>	Oct 29, 2012

GEL; TRANSDERMAL

DIVIGEL

<u>AB</u>	+! VERTICAL PHARMS	<u>0.1% (0.25GM/PACKET)</u>	<u>N022038 001</u>	Jun 04, 2007
<u>AB</u>	+!	<u>0.1% (0.5GM/PACKET)</u>	<u>N022038 002</u>	Jun 04, 2007
<u>AB</u>	+!	<u>0.1% (1GM/PACKET)</u>	<u>N022038 003</u>	Jun 04, 2007
<u>AB</u>	+!	<u>0.1% (0.75GM/PACKET)</u>	<u>N022038 004</u>	Aug 17, 2018
<u>AB</u>	+!	<u>0.1% (1.25GM/PACKET)</u>	<u>N022038 005</u>	Dec 12, 2019

PRESCRIPTION DRUG PRODUCT LIST

ESTRADIOL

GEL; TRANSDERMAL

ESTRADIOL

<u>AB</u>	AMNEAL	<u>0.1% (0.25GM/PACKET)</u>	<u>A216055 001</u>	Jul 12, 2024
<u>AB</u>		<u>0.1% (0.5GM/PACKET)</u>	<u>A216055 002</u>	Jul 12, 2024
<u>AB</u>		<u>0.1% (0.75GM/PACKET)</u>	<u>A216055 003</u>	Jul 12, 2024
<u>AB</u>		<u>0.1% (1GM/PACKET)</u>	<u>A216055 004</u>	Jul 12, 2024
<u>AB</u>		<u>0.1% (1.25GM/PACKET)</u>	<u>A216055 005</u>	Jul 12, 2024
<u>AB</u>	CHEMO RESEARCH SL	<u>0.1% (0.25GM/PACKET)</u>	<u>A211783 001</u>	Aug 10, 2022
<u>AB</u>		<u>0.1% (0.5GM/PACKET)</u>	<u>A211783 002</u>	Aug 10, 2022
<u>AB</u>		<u>0.1% (1GM/PACKET)</u>	<u>A211783 003</u>	Aug 10, 2022
<u>AB</u>	NOVITIUM PHARMA	<u>0.1% (0.25GM/PACKET)</u>	<u>A217610 001</u>	Aug 24, 2023
<u>AB</u>		<u>0.1% (0.5GM/PACKET)</u>	<u>A217610 002</u>	Aug 24, 2023
<u>AB</u>		<u>0.1% (0.75GM/PACKET)</u>	<u>A217610 003</u>	Aug 24, 2023
<u>AB</u>		<u>0.1% (1GM/PACKET)</u>	<u>A217610 004</u>	Aug 24, 2023
<u>AB</u>		<u>0.1% (1.25GM/PACKET)</u>	<u>A217610 005</u>	Aug 24, 2023
<u>AB</u>	PADAGIS ISRAEL	<u>0.1% (0.25GM/PACKET)</u>	<u>A216524 001</u>	Nov 14, 2023
<u>AB</u>		<u>0.1% (0.5GM/PACKET)</u>	<u>A216524 002</u>	Nov 14, 2023
<u>AB</u>		<u>0.1% (0.75GM/PACKET)</u>	<u>A216524 003</u>	Nov 14, 2023
<u>AB</u>		<u>0.1% (1GM/PACKET)</u>	<u>A216524 004</u>	Nov 14, 2023
<u>AB</u>		<u>0.1% (1.25GM/PACKET)</u>	<u>A216524 005</u>	Nov 14, 2023
<u>AB</u>	QUAGEN	<u>0.1% (0.25GM/PACKET)</u>	<u>A217863 001</u>	Jul 12, 2024
<u>AB</u>		<u>0.1% (0.5GM/PACKET)</u>	<u>A217863 002</u>	Jul 12, 2024
<u>AB</u>		<u>0.1% (0.75GM/PACKET)</u>	<u>A217863 003</u>	Jul 12, 2024
<u>AB</u>		<u>0.1% (1GM/PACKET)</u>	<u>A217863 004</u>	Jul 12, 2024
<u>AB</u>		<u>0.1% (1.25GM/PACKET)</u>	<u>A217863 005</u>	Jul 12, 2024

GEL, METERED; TRANSDERMAL

ESTRADIOL

<u>AB</u>	NOVITIUM PHARMA	<u>0.06% (1.25GM/ACTIVATION)</u>	<u>A217882 001</u>	Oct 23, 2024
<u>AB</u>	SOLARIS PHARMA CORP	<u>0.06% (1.25GM/ACTIVATION)</u>	<u>A216160 001</u>	Apr 22, 2024
<u>ESTROGEL</u>				
<u>AB</u>	+! ASCEND THERAPS US	<u>0.06% (1.25GM/ACTIVATION)</u>	<u>N021166 002</u>	Feb 09, 2004
ELESTRIN				
	+! MYLAN SPECIALITY LP	0.06% (0.87GM/ACTIVATION)	N021813 001	Dec 15, 2006
INSERT; VAGINAL				
IMVEXXY				
	+ MAYNE PHARMA	0.004MG	N208564 001	May 29, 2018
	+!	0.01MG	N208564 002	May 29, 2018
INSERT, EXTENDED RELEASE; VAGINAL				
ESTRING				
	+! PFIZER	0.0075MG/24HR	N020472 001	Apr 26, 1996
SPRAY; TRANSDERMAL				
EVAMIST				
	+! PADAGIS US	1.53MG/SPRAY	N022014 001	Jul 27, 2007

SYSTEM; TRANSDERMAL

ESTRADIOL

<u>AB</u>	ZYDUS PHARMS	<u>0.014MG/24HR</u>	<u>A204379 001</u>	Apr 17, 2023
<u>MENOSTAR</u>				
<u>AB</u>	+! BAYER HLTHCARE	<u>0.014MG/24HR</u>	<u>N021674 001</u>	Jun 08, 2004
<u>ESTRADIOL</u>				
<u>AB1</u>	AMNEAL	<u>0.025MG/24HR</u>	<u>A211293 001</u>	Feb 04, 2019
<u>AB1</u>		<u>0.0375MG/24HR</u>	<u>A211293 002</u>	Feb 04, 2019
<u>AB1</u>		<u>0.05MG/24HR</u>	<u>A211293 003</u>	Feb 04, 2019
<u>AB1</u>		<u>0.075MG/24HR</u>	<u>A211293 004</u>	Feb 04, 2019
<u>AB1</u>		<u>0.1MG/24HR</u>	<u>A211293 005</u>	Feb 04, 2019
<u>AB1</u>	MYLAN TECHNOLOGIES	<u>0.025MG/24HR</u>	<u>A201675 001</u>	Dec 19, 2014
<u>AB1</u>		<u>0.0375MG/24HR</u>	<u>A201675 002</u>	Dec 19, 2014
<u>AB1</u>		<u>0.05MG/24HR</u>	<u>A201675 003</u>	Dec 19, 2014
<u>AB1</u>		<u>0.075MG/24HR</u>	<u>A201675 004</u>	Dec 19, 2014
<u>AB1</u>		<u>0.1MG/24HR</u>	<u>A201675 005</u>	Dec 19, 2014
<u>AB1</u>	ZYDUS PHARMS	<u>0.025MG/24HR</u>	<u>A206241 001</u>	Dec 01, 2022
<u>AB1</u>		<u>0.0375MG/24HR</u>	<u>A206241 002</u>	Dec 01, 2022
<u>AB1</u>		<u>0.05MG/24HR</u>	<u>A206241 003</u>	Dec 01, 2022
<u>AB1</u>		<u>0.075MG/24HR</u>	<u>A206241 004</u>	Dec 01, 2022
<u>AB1</u>		<u>0.1MG/24HR</u>	<u>A206241 005</u>	Dec 01, 2022

VIVELLE-DOT

<u>AB1</u>	+ SANDOZ	<u>0.025MG/24HR</u>	<u>N020538 009</u>	May 03, 2002
<u>AB1</u>	+	<u>0.0375MG/24HR</u>	<u>N020538 005</u>	Jan 08, 1999
<u>AB1</u>	+	<u>0.05MG/24HR</u>	<u>N020538 006</u>	Jan 08, 1999
<u>AB1</u>	+	<u>0.075MG/24HR</u>	<u>N020538 007</u>	Jan 08, 1999
<u>AB1</u>	+!	<u>0.1MG/24HR</u>	<u>N020538 008</u>	Jan 08, 1999

PRESCRIPTION DRUG PRODUCT LIST

ESTRADIOL

TABLET; ORAL

ESTRADIOL

<u>AB</u>	BARR LABS INC	<u>0.5MG</u>	<u>A040197 001</u>	Oct 22, 1997
<u>AB</u>		<u>1MG</u>	<u>A040197 002</u>	Oct 22, 1997
<u>AB</u>	!	<u>2MG</u>	<u>A040197 003</u>	Oct 22, 1997
<u>AB</u>	EPIC PHARMA LLC	<u>0.5MG</u>	<u>A040275 001</u>	Dec 29, 1998
<u>AB</u>		<u>1MG</u>	<u>A040275 002</u>	Dec 29, 1998
<u>AB</u>		<u>2MG</u>	<u>A040275 003</u>	Dec 29, 1998
<u>AB</u>	NOVITIUM PHARMA	<u>0.5MG</u>	<u>A217334 001</u>	Sep 06, 2023
<u>AB</u>		<u>1MG</u>	<u>A217334 002</u>	Sep 06, 2023
<u>AB</u>		<u>2MG</u>	<u>A217334 003</u>	Sep 06, 2023

TABLET; VAGINAL

ESTRADIOL

<u>AB</u>	AMNEAL PHARMS	<u>10MCG</u>	<u>A205256 001</u>	May 29, 2015
<u>AB</u>	AUROBINDO PHARMA	<u>10MCG</u>	<u>A216550 001</u>	Aug 02, 2024
<u>AB</u>	GLENMARK PHARMS LTD	<u>10MCG</u>	<u>A210264 001</u>	Sep 14, 2018
<u>AB</u>	TEVA PHARMS USA	<u>10MCG</u>	<u>A206388 001</u>	Jul 21, 2017

VAGIFEM

<u>AB</u>	+!	NOVO NORDISK INC	<u>10MCG</u>	<u>N020908 002</u>	Nov 25, 2009
-----------	----	------------------	--------------	--------------------	--------------

ESTRADIOL ACETATE

INSERT, EXTENDED RELEASE; VAGINAL

FEMRING

+	MILLICENT PR	EQ 0.05MG BASE/24HR	N021367 001	Mar 20, 2003
+	!	EQ 0.1MG BASE/24HR	N021367 002	Mar 20, 2003

ESTRADIOL CYPIONATE

INJECTABLE; INJECTION

DEPO-ESTRADIOL

!	PFIZER	5MG/ML	A085470 003	
---	--------	--------	-------------	--

ESTRADIOL VALERATE

INJECTABLE; INJECTION

DELESTROGEN

<u>AO</u>	+!	ENDO OPERATIONS	<u>10MG/ML</u>	<u>N009402 002</u>	
<u>AO</u>	+!		<u>20MG/ML</u>	<u>N009402 004</u>	

ESTRADIOL VALERATE

<u>AO</u>		AM REGENT	<u>20MG/ML</u>	<u>A090920 001</u>	Jan 19, 2010
<u>AO</u>			<u>40MG/ML</u>	<u>A090920 002</u>	Jan 19, 2010
<u>AO</u>		HIKMA	<u>10MG/ML</u>	<u>A203723 001</u>	Apr 21, 2020
<u>AO</u>			<u>20MG/ML</u>	<u>A203723 002</u>	Apr 21, 2020
<u>AO</u>			<u>40MG/ML</u>	<u>A203723 003</u>	Apr 21, 2020
<u>AO</u>		XIROMED	<u>10MG/ML</u>	<u>A216656 001</u>	Apr 28, 2023
<u>AO</u>			<u>20MG/ML</u>	<u>A216656 002</u>	Apr 28, 2023
<u>AO</u>	!		<u>40MG/ML</u>	<u>A216656 003</u>	Apr 28, 2023

ESTRADIOL; LEVONORGESTREL

FILM, EXTENDED RELEASE; TRANSDERMAL

CLIMARA PRO

+	!	BAYER HLTHCARE	0.045MG/24HR; 0.015MG/24HR	N021258 001	Nov 21, 2003
---	---	----------------	----------------------------	-------------	--------------

ESTRADIOL; NORETHINDRONE ACETATE

FILM, EXTENDED RELEASE; TRANSDERMAL

COMBIPATCH

+		NOVEN PHARMS INC	0.05MG/24HR; 0.14MG/24HR	N020870 001	Aug 07, 1998
+	!		0.05MG/24HR; 0.25MG/24HR	N020870 002	Aug 07, 1998

TABLET; ORAL

ACTIVELLA

<u>AB</u>	+!	AMNEAL	<u>1MG; 0.5MG</u>	<u>N020907 001</u>	Nov 18, 1998
-----------	----	--------	-------------------	--------------------	--------------

ESTRADIOL AND NORETHINDRONE ACETATE

<u>AB</u>		BARR	<u>1MG; 0.5MG</u>	<u>A079193 001</u>	May 11, 2010
<u>AB</u>		BRECKENRIDGE PHARM	<u>0.5MG; 0.1MG</u>	<u>A078324 002</u>	Jun 09, 2011
<u>AB</u>			<u>1MG; 0.5MG</u>	<u>A078324 001</u>	Apr 17, 2008
<u>AB</u>		NAARI PTE LTD	<u>1MG; 0.5MG</u>	<u>A210233 001</u>	Feb 28, 2018
<u>AB</u>		NOVAST LABS	<u>0.5MG; 0.1MG</u>	<u>A210612 001</u>	Apr 03, 2019
<u>AB</u>			<u>1MG; 0.5MG</u>	<u>A210612 002</u>	Apr 03, 2019
<u>AB</u>		XIROMED	<u>0.5MG; 0.1MG</u>	<u>A207261 001</u>	Feb 10, 2017
<u>AB</u>			<u>1MG; 0.5MG</u>	<u>A207261 002</u>	Feb 10, 2017

AMABELZ

BX		LUPIN LTD	0.5MG; 0.1MG	A203339 001	Jun 20, 2016
BX			1MG; 0.5MG	A203339 002	Jun 20, 2016

PRESCRIPTION DRUG PRODUCT LISTESTRADIOL; NORETHINDRONE ACETATE; RELUGOLIX

TABLET; ORAL

MYFEMBREE

+! SUMITOMO PHARMA 1MG; 0.5MG; 40MG N214846 001 May 26, 2021

ESTRADIOL; PROGESTERONE

CAPSULE; ORAL

BIJUVA

+ MAYNE PHARMA 0.5MG; 100MG N210132 002 Dec 28, 2021

+! 1MG; 100MG N210132 001 Oct 28, 2018

ESTROGENS, CONJUGATED

CREAM; TOPICAL, VAGINAL

PREMARIN

+! WYETH PHARMS 0.625MG/GM N020216 001

INJECTABLE; INJECTION

PREMARIN

+! WYETH PHARMS 25MG/VIAL N010402 001

TABLET; ORAL

PREMARIN

+ WYETH PHARMS 0.3MG N004782 003

+ 0.45MG N004782 006 Jul 16, 2003

+! 0.625MG N004782 004

+! 0.9MG N004782 005 Jan 26, 1984

+! 1.25MG N004782 001

ESTROGENS, CONJUGATED; MEDROXYPROGESTERONE ACETATE

TABLET; ORAL-28

PREMPHASE 14/14

+! WYETH PHARMS 0.625MG, 0.625MG; N/A, 5MG N020527 002 Nov 17, 1995

PREMPRO

+! WYETH PHARMS 0.3MG; 1.5MG N020527 005 Jun 04, 2003

+! 0.45MG; 1.5MG N020527 004 Mar 12, 2003

+! 0.625MG; 2.5MG N020527 001 Nov 17, 1995

+! 0.625MG; 5MG N020527 003 Jan 09, 1998

ESTROPIPATE

TABLET; ORAL

OGEN 5

+ PFIZER 6MG A083220 004

ESZOPICLONE

TABLET; ORAL

ESZOPICLONE

<u>AB</u>	AUROBINDO PHARMA	<u>1MG</u>	<u>A208451 001</u>	Sep 15, 2016
<u>AB</u>		<u>2MG</u>	<u>A208451 002</u>	Sep 15, 2016
<u>AB</u>		<u>3MG</u>	<u>A208451 003</u>	Sep 15, 2016
<u>AB</u>	DR REDDYS	<u>1MG</u>	<u>A091024 001</u>	Apr 15, 2014
<u>AB</u>		<u>2MG</u>	<u>A091024 002</u>	Apr 15, 2014
<u>AB</u>		<u>3MG</u>	<u>A091024 003</u>	Apr 15, 2014
<u>AB</u>	GLENMARK PHARMS LTD	<u>1MG</u>	<u>A091166 001</u>	Apr 15, 2014
<u>AB</u>		<u>2MG</u>	<u>A091166 002</u>	Apr 15, 2014
<u>AB</u>		<u>3MG</u>	<u>A091166 003</u>	Apr 15, 2014
<u>AB</u>	HETERO LABS LTD V	<u>1MG</u>	<u>A205504 001</u>	Jan 04, 2024
<u>AB</u>		<u>2MG</u>	<u>A205504 002</u>	Jan 04, 2024
<u>AB</u>		<u>3MG</u>	<u>A205504 003</u>	Jan 04, 2024
<u>AB</u>	LUPIN LTD	<u>1MG</u>	<u>A091124 001</u>	Sep 13, 2011
<u>AB</u>		<u>2MG</u>	<u>A091124 002</u>	Sep 13, 2011
<u>AB</u>		<u>3MG</u>	<u>A091124 003</u>	Sep 13, 2011
<u>AB</u>	MACLEODS PHARMS LTD	<u>1MG</u>	<u>A202929 001</u>	Jan 30, 2015
<u>AB</u>		<u>2MG</u>	<u>A202929 002</u>	Jan 30, 2015
<u>AB</u>		<u>3MG</u>	<u>A202929 003</u>	Jan 30, 2015
<u>AB</u>	MYLAN PHARMS INC	<u>1MG</u>	<u>A091151 001</u>	Mar 26, 2013
<u>AB</u>		<u>2MG</u>	<u>A091151 002</u>	Mar 26, 2013
<u>AB</u>		<u>3MG</u>	<u>A091151 003</u>	Mar 26, 2013
<u>AB</u>	ORBION PHARMS	<u>1MG</u>	<u>A091113 001</u>	Jun 10, 2014
<u>AB</u>		<u>2MG</u>	<u>A091113 002</u>	Jun 10, 2014
<u>AB</u>		<u>3MG</u>	<u>A091113 003</u>	Jun 10, 2014
<u>AB</u>	SUN PHARM	<u>1MG</u>	<u>A091103 001</u>	Apr 03, 2013
<u>AB</u>		<u>2MG</u>	<u>A091103 002</u>	Apr 03, 2013
<u>AB</u>		<u>3MG</u>	<u>A091103 003</u>	Apr 03, 2013
<u>AB</u>	TEVA	<u>1MG</u>	<u>A091169 001</u>	May 23, 2011
<u>AB</u>		<u>2MG</u>	<u>A091169 002</u>	May 23, 2011
<u>AB</u>		<u>3MG</u>	<u>A091169 003</u>	May 23, 2011

PRESCRIPTION DRUG PRODUCT LIST

ESZOPICLONE

TABLET; ORAL

LUNESTA

AB	+	WAYLIS THERAP	1MG	N021476 001	Dec 15, 2004
AB	+		2MG	N021476 002	Dec 15, 2004
AB	+	!	3MG	N021476 003	Dec 15, 2004

ETELICALCETIDE

SOLUTION; INTRAVENOUS

PARSABIV

+	!	KAI PHARMS INC	2.5MG/0.5ML (2.5MG/0.5ML)	N208325 001	Feb 07, 2017
+	!		5MG/ML (5MG/ML)	N208325 002	Feb 07, 2017
+	!		10MG/2ML (5MG/ML)	N208325 003	Feb 07, 2017

ETEPLIRSEN

SOLUTION; INTRAVENOUS

EXONDYS 51

+	!	SAREPTA THERAPS INC	100MG/2ML (50MG/ML)	N206488 001	Sep 19, 2016
+	!		500MG/10ML (50MG/ML)	N206488 002	Sep 19, 2016

ETHACRYNATE SODIUM

INJECTABLE; INJECTION

EDECRI

AP	+	!	BAUSCH	EQ 50MG BASE/VIAL	N016093 001
-----------	---	---	--------	--------------------------	--------------------

ETHACRYNATE SODIUM

AP		ENDO OPERATIONS	EQ 50MG BASE/VIAL	A205473 001	Jul 29, 2015
AP		MYLAN INSTITUTIONAL	EQ 50MG BASE/VIAL	A204634 001	Aug 23, 2016
AP		STERIMAX	EQ 50MG BASE/VIAL	A208663 001	Jun 09, 2020
AP		ZYDUS PHARMS	EQ 50MG BASE/VIAL	A207758 001	Nov 17, 2017

ETHACRYNIC ACID

TABLET; ORAL

EDECRI

AB	+	!	BAUSCH	25MG	N016092 001
-----------	---	---	--------	-------------	--------------------

ETHACRYNIC ACID

AB		ADAPTIS	25MG	A205609 001	Jun 30, 2016
AB		AGNITIO	25MG	A211809 001	Jul 12, 2019
AB		AMNEAL PHARMS CO	25MG	A208805 001	May 08, 2018
AB		CHARTWELL RX	25MG	A213240 001	Oct 19, 2020
AB		LUPIN LTD	25MG	A211719 001	Sep 06, 2019
AB		SCIEGEN PHARMS INC	25MG	A211232 001	Aug 27, 2019
AB		UPSHER SMITH LABS	25MG	A212417 001	Feb 19, 2020

ETHAMBUTOL HYDROCHLORIDE

TABLET; ORAL

ETHAMBUTOL HYDROCHLORIDE

AB		EPIC PHARMA LLC	100MG	A075095 001	Nov 30, 1999
AB			400MG	A075095 002	Nov 30, 1999
AB		LUPIN	100MG	A078939 001	Jun 17, 2009
AB			400MG	A078939 002	Jun 17, 2009
AB	+	KANCHAN HLTHCARE	100MG	N016320 001	
AB	+	!	400MG	N016320 003	

ETHANOLAMINE OLEATE

INJECTABLE; INJECTION

ETHAMOLIN

+	!	QOL MEDCL	50MG/ML	N019357 001	Dec 22, 1988
---	---	-----------	---------	-------------	--------------

ETHINYL ESTRADIOL; ETHYNODIOL DIACETATE

TABLET; ORAL-28

ETHYNODIOL DIACETATE AND ETHINYL ESTRADIOL

AB		XIROMED	0.035MG;1MG	A204703 001	Jul 28, 2016
AB			0.05MG;1MG	A204704 001	Feb 09, 2016
AB		KELNOR			
AB		BARR	0.035MG;1MG	A076785 001	May 23, 2005
AB		LO-MALMOREDE			
AB		NOVAST LABS	0.035MG;1MG	A209548 001	Feb 11, 2019
AB		MALMOREDE			
AB		NOVAST LABS	0.05MG;1MG	A209547 001	Jul 25, 2018
AB		ZOVIA 1/50E-28			
AB	!	WATSON LABS	0.05MG;1MG	A072723 001	Dec 30, 1991

PRESCRIPTION DRUG PRODUCT LIST

ETHINYL ESTRADIOL; ETONOGESTREL

RING; VAGINAL

ELURYNG

AB	AMNEAL	<u>0.015MG/24HR;0.12MG/24HR</u>	<u>A210830</u>	<u>001</u>	Dec 11, 2019
-----------	--------	---------------------------------	----------------	------------	--------------

ENILLORING

AB	XIROMED	<u>0.015MG/24HR;0.12MG/24HR</u>	<u>A211157</u>	<u>001</u>	Jun 29, 2023
-----------	---------	---------------------------------	----------------	------------	--------------

ETHINYL ESTRADIOL; ETONOGESTREL

AB	TEVA PHARMS USA INC	<u>0.015MG/24HR;0.12MG/24HR</u>	<u>A204305</u>	<u>001</u>	Jan 13, 2021
-----------	---------------------	---------------------------------	----------------	------------	--------------

HALOETTE

AB	DR REDDYS LABS SA	<u>0.015MG/24HR;0.12MG/24HR</u>	<u>A211328</u>	<u>001</u>	Aug 05, 2022
-----------	-------------------	---------------------------------	----------------	------------	--------------

NUVARING

AB	+! ORGANON USA ORGANON	<u>0.015MG/24HR;0.12MG/24HR</u>	<u>N021187</u>	<u>001</u>	Oct 03, 2001
-----------	------------------------	---------------------------------	----------------	------------	--------------

ETHINYL ESTRADIOL; LEVONORGESTREL

SYSTEM; TRANSDERMAL

TWIRLA

+! AGILE

0.03MG/24HR;0.12MG/24HR

N204017 001 Feb 14, 2020

TABLET; ORAL

ASHLYNA

AB	GLENMARK PHARMS LTD	<u>0.03MG,0.01MG;0.15MG,N/A</u>	<u>A203163</u>	<u>001</u>	Feb 23, 2015
-----------	---------------------	---------------------------------	----------------	------------	--------------

DAYSEE

AB	LUPIN LTD	<u>0.03MG,0.01MG;0.15MG,N/A</u>	<u>A091467</u>	<u>001</u>	Apr 10, 2013
-----------	-----------	---------------------------------	----------------	------------	--------------

DOLISHALE

AB	NOVAST LABS	<u>0.02MG;0.09MG</u>	<u>A091692</u>	<u>001</u>	Oct 22, 2020
-----------	-------------	----------------------	----------------	------------	--------------

ICLEVIA

AB	AUROBINDO PHARMA LTD	<u>0.03MG;0.15MG</u>	<u>A206850</u>	<u>001</u>	Jun 29, 2018
-----------	----------------------	----------------------	----------------	------------	--------------

INTROVALE

AB	XIROMED	<u>0.03MG;0.15MG</u>	<u>A079064</u>	<u>001</u>	Sep 27, 2010
-----------	---------	----------------------	----------------	------------	--------------

JAIMIESS

AB	XIROMED	<u>0.03MG,0.01MG;0.15MG,N/A</u>	<u>A203770</u>	<u>001</u>	Dec 27, 2017
-----------	---------	---------------------------------	----------------	------------	--------------

LEVONORGESTREL AND ETHINYL ESTRADIOL

AB	AMNEAL PHARMS	<u>0.03MG;0.15MG</u>	<u>A203871</u>	<u>001</u>	Nov 13, 2015
-----------	---------------	----------------------	----------------	------------	--------------

AB		<u>0.03MG,0.01MG;0.15MG,N/A</u>	<u>A203872</u>	<u>001</u>	Dec 22, 2015
-----------	--	---------------------------------	----------------	------------	--------------

AB	GLENMARK PHARMS LTD	<u>0.02MG;0.09MG</u>	<u>A202791</u>	<u>001</u>	Apr 09, 2015
-----------	---------------------	----------------------	----------------	------------	--------------

AB		<u>0.03MG;0.15MG</u>	<u>A203164</u>	<u>001</u>	Jun 12, 2015
-----------	--	----------------------	----------------	------------	--------------

AB	LUPIN LTD	<u>0.03MG;0.15MG</u>	<u>A091440</u>	<u>001</u>	Oct 23, 2012
-----------	-----------	----------------------	----------------	------------	--------------

AB	! WATSON LABS	<u>0.02MG;0.09MG</u>	<u>A079218</u>	<u>001</u>	Jun 06, 2011
-----------	---------------	----------------------	----------------	------------	--------------

AB	XIROMED	<u>0.03MG;0.15MG</u>	<u>A200490</u>	<u>001</u>	Apr 21, 2015
-----------	---------	----------------------	----------------	------------	--------------

LEVONORGESTREL AND ETHINYL ESTRADIOL AND ETHINYL ESTRADIOL

AB	DR REDDYS LABS SA	<u>0.03MG,0.01MG;0.15MG,N/A</u>	<u>A078834</u>	<u>001</u>	May 31, 2011
-----------	-------------------	---------------------------------	----------------	------------	--------------

AB	LUPIN LTD	<u>0.02MG,0.1MG;0.01MG,N/A</u>	<u>A091674</u>	<u>001</u>	Oct 26, 2011
-----------	-----------	--------------------------------	----------------	------------	--------------

AB	XIROMED	<u>0.02MG,0.15MG;0.025MG,0.15MG;0.03MG,0.15MG;0.01MG,N/A</u>	<u>A206053</u>	<u>001</u>	Oct 02, 2017
-----------	---------	--	----------------	------------	--------------

AB		<u>0.02MG,0.1MG;0.01MG,N/A</u>	<u>A200493</u>	<u>001</u>	Jun 17, 2015
-----------	--	--------------------------------	----------------	------------	--------------

AB		<u>0.02MG,0.1MG;0.01MG,N/A</u>	<u>A205131</u>	<u>001</u>	Dec 14, 2017
-----------	--	--------------------------------	----------------	------------	--------------

AB		<u>0.03MG,0.01MG;0.15MG,N/A</u>	<u>A200492</u>	<u>001</u>	May 27, 2015
-----------	--	---------------------------------	----------------	------------	--------------

LO SIMPESS

AB	AUROBINDO PHARMA	<u>0.02MG,0.1MG;0.01MG,N/A</u>	<u>A206852</u>	<u>001</u>	Apr 28, 2017
-----------	------------------	--------------------------------	----------------	------------	--------------

LOSEASONIQUE

AB	TEVA BRANDED PHARM	<u>0.02MG,0.1MG;0.01MG,N/A</u>	<u>N022262</u>	<u>001</u>	Oct 24, 2008
-----------	--------------------	--------------------------------	----------------	------------	--------------

QUARTETTE

AB	+! TEVA BRANDED PHARM	<u>0.02MG,0.15MG;0.025MG,0.15MG;0.03MG,0.15MG;0.01MG,N/A</u>	<u>N204061</u>	<u>001</u>	Mar 28, 2013
-----------	-----------------------	--	----------------	------------	--------------

QUASENSE

AB	WATSON LABS	<u>0.03MG;0.15MG</u>	<u>A077101</u>	<u>001</u>	Sep 06, 2006
-----------	-------------	----------------------	----------------	------------	--------------

SEASONALE

AB	+! TEVA BRANDED PHARM	<u>0.03MG;0.15MG</u>	<u>N021544</u>	<u>001</u>	Sep 05, 2003
-----------	-----------------------	----------------------	----------------	------------	--------------

SEASONIQUE

AB	+! TEVA BRANDED PHARM	<u>0.03MG,0.01MG;0.15MG,N/A</u>	<u>N021840</u>	<u>001</u>	May 25, 2006
-----------	-----------------------	---------------------------------	----------------	------------	--------------

SETLAKIN

AB	NOVAST LABS	<u>0.03MG;0.15MG</u>	<u>A090716</u>	<u>001</u>	Sep 15, 2014
-----------	-------------	----------------------	----------------	------------	--------------

SIMPESSE

AB	AUROBINDO PHARMA	<u>0.03MG,0.01MG;0.15MG,N/A</u>	<u>A206851</u>	<u>001</u>	Apr 07, 2017
-----------	------------------	---------------------------------	----------------	------------	--------------

BALCOLTRA

AB3	+! AVION PHARMS	<u>0.02MG;0.1MG</u>	<u>N208612</u>	<u>001</u>	Jan 09, 2018
------------	-----------------	---------------------	----------------	------------	--------------

LEVONORGESTREL AND ETHINYL ESTRADIOL AND FERROUS FUMARATE

AB3	XIROMED	<u>0.02MG;0.1MG</u>	<u>A214640</u>	<u>001</u>	Aug 16, 2023
------------	---------	---------------------	----------------	------------	--------------

MINZOYA

AB3	LUPIN LTD	<u>0.02MG;0.1MG</u>	<u>A217087</u>	<u>001</u>	Feb 15, 2024
------------	-----------	---------------------	----------------	------------	--------------

TYBLUME

+! EXELTIS USA INC

0.02MG;0.1MG

N209405 001 Mar 30, 2020

PRESCRIPTION DRUG PRODUCT LIST

ETHINYL ESTRADIOL; LEVONORGESTREL

TABLET; ORAL-28

<u>ALTAVERA</u>					
AB	XIROMED	0.03MG;0.15MG	A079102	001	Aug 03, 2010
<u>AYUNA</u>					
AB	AUROBINDO PHARMA	0.03MG;0.15MG	A206866	001	Sep 23, 2016
<u>ENPRESSE-28</u>					
AB	DURAMED PHARMS BARR	0.03MG,0.04MG,0.03MG;0.05MG,0.075MG,0.125MG	A075809	002	Jul 16, 2001
<u>KURVELO</u>					
AB	LUPIN LTD	0.03MG;0.15MG	A091408	001	Oct 17, 2012
<u>LEVONEST</u>					
AB	NOVAST LABS LTD	0.03MG,0.04MG,0.03MG;0.05MG,0.075MG,0.125MG	A090719	001	Dec 29, 2010
<u>LEVONORGESTREL AND ETHINYL ESTRADIOL</u>					
AB	LUPIN LTD	0.03MG,0.04MG,0.03MG;0.05MG,0.075MG,0.125MG	A200248	001	Nov 19, 2015
AB	NAARI PTE LTD	0.03MG;0.15MG	A207033	001	Oct 09, 2020
AB	XIROMED	0.03MG;0.15MG	A091663	001	Dec 21, 2012
<u>LEVORA 0.15/30-28</u>					
AB	! DR REDDYS LABS SA	0.03MG;0.15MG	A073594	001	Dec 13, 1993
<u>MARLISSA</u>					
AB	GLENMARK PHARMS LTD	0.03MG;0.15MG	A091452	001	Feb 29, 2012
<u>MYZILRA</u>					
AB	ENDO OPERATIONS	0.03MG,0.04MG,0.03MG;0.05MG,0.075MG,0.125MG	A077502	001	Nov 23, 2011
<u>PORTIA-28</u>					
AB	BARR	0.03MG;0.15MG	A075866	002	May 23, 2002
<u>TRIVORA-28</u>					
AB	! DR REDDYS LABS SA	0.03MG,0.04MG,0.03MG;0.05MG,0.075MG,0.125MG	A074538	002	Dec 18, 1997
<u>AFIRMELLE</u>					
AB1	AUROBINDO PHARMA	0.02MG;0.1MG	A206886	001	Nov 14, 2016
<u>AVIANE-28</u>					
AB1	DURAMED PHARMS BARR	0.02MG;0.1MG	A075796	001	Apr 30, 2001
<u>FALMINA</u>					
AB1	NOVAST LABS LTD	0.02MG;0.1MG	A090721	001	Mar 28, 2012
<u>LEVONORGESTREL AND ETHINYL ESTRADIOL</u>					
AB1	! DR REDDYS LABS SA	0.02MG;0.1MG	A076625	001	Nov 18, 2004
AB1	HETERO LABS	0.02MG;0.1MG	A212298	001	Feb 13, 2023
AB1	LUPIN LTD	0.02MG;0.1MG	A091425	001	Jan 18, 2013
AB1	NAARI PTE LTD	0.02MG;0.1MG	A207065	001	Aug 17, 2020
AB1	XIROMED	0.02MG;0.1MG	A200245	001	Oct 09, 2013
<u>VIENVA</u>					
AB1	XIROMED	0.02MG;0.1MG	A201088	001	May 21, 2015
<u>LESSINA-28</u>					
AB2	BARR	0.02MG;0.1MG	A075803	002	Mar 20, 2002
<u>LEVONORGESTREL AND ETHINYL ESTRADIOL</u>					
AB2	! DR REDDYS LABS SA	0.02MG;0.1MG	A077681	001	May 31, 2006

ETHINYL ESTRADIOL; NORELGESTROMIN

FILM, EXTENDED RELEASE; TRANSDERMAL

<u>ETHINYL ESTRADIOL AND NORELGESTROMIN</u>					
AB	AMNEAL	0.035MG/24HR;0.15MG/24HR	A213950	001	Feb 25, 2021
AB	ZYDUS PHARMS	0.035MG/24HR;0.15MG/24HR	A214594	001	Sep 14, 2023
<u>ONSURA</u>					
AB	TEVA PHARMS USA	0.035MG/24HR;0.15MG/24HR	A213977	001	Aug 25, 2021
<u>XULANE</u>					
AB	! MYLAN TECHNOLOGIES	0.035MG/24HR;0.15MG/24HR	A200910	001	Apr 16, 2014

ETHINYL ESTRADIOL; NORETHINDRONE

TABLET; ORAL

<u>NEXESTA FE</u>					
AB	AUROBINDO PHARMA	0.035MG;0.4MG	A207535	001	Feb 02, 2017
<u>NORETHINDRONE AND ETHINYL ESTRADIOL AND FERROUS FUMARATE</u>					
AB	AMNEAL PHARMS	0.035MG;0.4MG	A078892	001	Sep 26, 2011
AB	BARR	0.035MG;0.4MG	A078965	001	Aug 05, 2010
AB	LUPIN LTD	0.035MG;0.4MG	A091332	001	Mar 23, 2016
AB	NAARI PTE LTD	0.035MG;0.4MG	A207066	001	Mar 29, 2017
AB	! XIROMED	0.035MG;0.4MG	A202086	001	Apr 01, 2015
TABLET; ORAL-21					
<u>NORTREL 1/35-21</u>					
AB	BARR	0.035MG;1MG	A072693	001	Feb 28, 1992

PRESCRIPTION DRUG PRODUCT LIST

ETHINYL ESTRADIOL; NORETHINDRONE

TABLET; ORAL-21

NORTREL 7/7/7

BARR

0.035MG, 0.035MG, 0.035MG; 0.5MG, 0.75MG, 1MG

A075478 001 Aug 30, 2002

TABLET; ORAL-28

ALYACEN 1/35

AB GLENMARK PHARMS LTD

0.035MG; 1MGA091634 001 Jan 19, 2012ALYACEN 7/7/7

AB GLENMARK PHARMS LTD

0.035MG, 0.035MG, 0.035MG; 0.5MG, 0.75MG, 1MGA091636 001 Jan 19, 2012ARANELLE

AB BARR

0.035MG, 0.035MG, 0.035MG; 0.5MG, 1MG, 0.5MGA076783 001 Sep 29, 2004BALZIVA-28

AB ! BARR

0.035MG; 0.4MGA076238 001 Apr 22, 2004BRIELLYN

AB GLENMARK PHARMS LTD

0.035MG; 0.4MGA090538 001 Mar 22, 2011CYONANZ

AB AUROBINDO PHARMA

0.035MG; 0.5MGA207055 001 Oct 21, 2016DASETTA 1/35

AB NOVAST LABS LTD

0.035MG; 1MGA090948 001 Dec 22, 2011DASETTA 7/7/7

AB ! NOVAST LABS LTD

0.035MG, 0.035MG, 0.035MG; 0.5MG, 0.75MG, 1MGA090946 001 Dec 22, 2011GILDAGIA

AB ENDO OPERATIONS

0.035MG; 0.4MGA078376 001 Nov 06, 2012NORETHINDRONE AND ETHINYL ESTRADIOL

AB NAARI PTE LTD

0.035MG; 1MGA206864 001 Apr 28, 2017

AB WATSON LABS

0.035MG; 0.4MGA078323 001 Feb 04, 2010

AB WATSON LABS TEVA

0.035MG; 1MGA070687 001 Jan 29, 1987NORTREL 0.5/35-28

AB BARR

0.035MG; 0.5MGA072695 001 Feb 28, 1992NORTREL 1/35-28

AB ! BARR

0.035MG; 1MGA072696 001 Feb 28, 1992NORTREL 7/7/7

AB BARR

0.035MG, 0.035MG, 0.035MG; 0.5MG, 0.75MG, 1MGA075478 002 Aug 30, 2002NYLIA 1/35

AB AUROBINDO PHARMA

0.035MG; 1MGA207056 001 Oct 21, 2016NYLIA 7/7/7

AB AUROBINDO PHARMA

0.035MG, 0.035MG, 0.035MG; 0.5MG, 0.75MG, 1MGA207054 001 Oct 21, 2016PHILITH

AB NOVAST LABS LTD

0.035MG; 0.4MGA090947 001 Dec 22, 2011TRI-NORINYL 28-DAY

AB +! DR REDDYS LABS SA

0.035MG, 0.035MG, 0.035MG; 0.5MG, 1MG, 0.5MGN018977 002 Apr 13, 1984VYFEMLA

AB LUPIN LTD

0.035MG; 0.4MGA201886 001 Sep 26, 2013WERA

AB ! NOVAST LABS LTD

0.035MG; 0.5MGA091204 001 Mar 27, 2012

NORETHINDRONE AND ETHINYL ESTRADIOL (10/11)

WATSON LABS TEVA

0.035MG, 0.035MG; 0.5MG, 1MG

A071044 001 Apr 01, 1988

TABLET, CHEWABLE; ORAL

KAITLIB FE

AB LUPIN LTD

0.025MG; 0.8MGA203448 001 Dec 17, 2015NORETHINDRONE AND ETHINYL ESTRADIOL AND FERROUS FUMARATE

AB +! TEVA BRANDED PHARM

0.025MG; 0.8MGN022573 001 Dec 22, 2010

AB XIROMED

0.025MG; 0.8MGA203371 001 Apr 23, 2014ETHINYL ESTRADIOL; NORETHINDRONE ACETATE

CAPSULE; ORAL

GEMMILY

AB XIROMED

0.02MG; 1MGA213317 001 Nov 09, 2020MERZEE

AB WILSHIRE PHARMS INC

0.02MG; 1MGA212706 001 Dec 18, 2020NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL AND FERROUS FUMARATE

AB GLENMARK PHARMS LTD

0.02MG; 1MGA213418 001 Jul 27, 2022TAYTULLA

AB +! APIL

0.02MG; 1MGN204426 001 Apr 19, 2013

TABLET; ORAL

AUROVELA 24 FE

AB AUROBINDO PHARMA

0.02MG; 1MGA207504 001 Jun 15, 2017

PRESCRIPTION DRUG PRODUCT LIST

ETHINYL ESTRADIOL; NORETHINDRONE ACETATE

TABLET; ORAL

<u>BLISOVI 24 FE</u>					
AB	LUPIN LTD	0.02MG;1MG	A091398	001	Oct 28, 2015
<u>FINZALA</u>					
AB	TEVA PHARMS USA INC	0.02MG;1MG	A210087	001	Apr 07, 2020
<u>FYAVOLV</u>					
AB	LUPIN LTD	0.005MG;1MG	A204213	002	Dec 10, 2015
AB		0.0025MG;0.5MG	A204213	001	Dec 10, 2015
<u>GILDESS 24 FE</u>					
AB	ENDO OPERATIONS	0.02MG;1MG	A090293	001	Dec 01, 2014
<u>HAILEY 24 FE</u>					
AB	! GLENMARK PHARMS LTD	0.02MG;1MG	A204847	001	Nov 17, 2017
<u>LARIN 24 FE</u>					
AB	NOVAST LABS	0.02MG;1MG	A202994	001	Feb 18, 2015
<u>LERIBANE</u>					
AB	NOVAST LABS	0.0025MG;0.5MG	A203435	002	Jun 03, 2016
AB		0.005MG;1MG	A203435	001	Jun 03, 2016
<u>MIBELAS 24 FE</u>					
AB	LUPIN	0.02MG;1MG	A206287	001	May 24, 2016
<u>NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL</u>					
AB	! BARR LABS INC	0.005MG;1MG	A076221	001	Nov 06, 2009
AB	GLENMARK PHARMS LTD	0.0025MG;0.5MG	A203038	001	Apr 02, 2015
AB		0.005MG;1MG	A203038	002	Apr 02, 2015
<u>NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL AND FERROUS FUMARATE</u>					
AB	BARR LABS INC	0.02MG;1MG	A090938	001	Dec 01, 2014
AB	! GLENMARK PHARMS LTD	0.02MG;1MG	A210369	001	Dec 26, 2017
AB	XIROMED	0.02MG;1MG	A209609	001	Jul 16, 2018
LO LOESTRIN FE					
	+! APIL	0.01MG,0.01MG;1MG,N/A	N022501	001	Oct 21, 2010
TABLET; ORAL-21					
<u>AUROVELA 1.5/30</u>					
AB	AUROBINDO PHARMA	0.03MG;1.5MG	A207581	001	Jun 26, 2017
<u>AUROVELA 1/20</u>					
AB	AUROBINDO PHARMA	0.02MG;1MG	A207506	001	Jun 16, 2017
<u>HAILEY 1.5/30</u>					
AB	GLENMARK SPECLT	0.03MG;1.5MG	A209297	001	Jun 05, 2018
<u>JUNEL 1.5/30</u>					
AB	BARR	0.03MG;1.5MG	A076381	001	May 30, 2003
<u>JUNEL 1/20</u>					
AB	BARR	0.02MG;1MG	A076380	001	May 30, 2003
<u>LARIN 1.5/30</u>					
AB	NOVAST LABS	0.03MG;1.5MG	A202996	001	Mar 20, 2014
<u>LARIN 1/20</u>					
AB	NOVAST LABS	0.02MG;1MG	A202995	001	Dec 04, 2013
<u>NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL</u>					
AB	GLENMARK PHARMS LTD	0.02MG;1MG	A206969	001	Jan 20, 2016
AB	XIROMED	0.02MG;1MG	A202771	001	Nov 06, 2013
AB		0.03MG;1.5MG	A202770	001	Feb 19, 2015
TABLET; ORAL-28					
<u>AUROVELA FE 1.5/30</u>					
AB	AUROBINDO PHARMA	0.03MG;1.5MG	A207580	001	Jun 15, 2017
<u>AUROVELA FE 1/20</u>					
AB	AUROBINDO PHARMA	0.02MG;1MG	A207505	001	Jun 16, 2017
<u>BLISOVI FE 1/20</u>					
AB	LUPIN LTD	0.02MG;1MG	A201584	001	Nov 18, 2015
<u>CHABELINA FE</u>					
AB	! NOVAST LABS	0.02MG,0.03MG,0.035MG;1MG,1MG,1MG	A202962	001	Apr 15, 2020
<u>HAILEY FE 1.5/30</u>					
AB	GLENMARK SPECLT	0.03MG;1.5MG	A209031	001	Jun 05, 2018
<u>HAILEY FE 1/20</u>					
AB	GLENMARK PHARMS LTD	0.02MG;1MG	A206597	001	Nov 21, 2017
<u>JUNEL FE 1.5/30</u>					
AB	! BARR	0.03MG;1.5MG	A076064	001	Sep 18, 2003
<u>JUNEL FE 1/20</u>					
AB	BARR	0.02MG;1MG	A076081	001	Sep 18, 2003
<u>LARIN FE 1.5/30</u>					
AB	NOVAST LABS	0.03MG;1.5MG	A091453	001	Aug 23, 2013
<u>LARIN FE 1/20</u>					
AB	NOVAST LABS	0.02MG;1MG	A091454	001	Aug 26, 2013
<u>NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL</u>					
AB	XIROMED	0.02MG;1MG	A202772	001	Nov 14, 2013

PRESCRIPTION DRUG PRODUCT LIST

ETHINYL ESTRADIOL; NORETHINDRONE ACETATE

TABLET; ORAL-28

NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL

<u>AB</u>		<u>0.02MG, 0.03MG, 0.035MG; 1MG, 1MG, 1MG</u>	<u>A205069 001</u>	Jun 22, 2018
<u>AB</u>		<u>0.03MG; 1.5MG</u>	<u>A202741 001</u>	Feb 20, 2015
	<u>TRI-LEGEST FE</u>			
<u>AB</u>	BARR	<u>0.02MG, 0.03MG, 0.035MG; 1MG, 1MG, 1MG</u>	<u>A076105 001</u>	Oct 26, 2007
	BLISOVI FE 1.5/30			
<u>BX</u>	LUPIN LTD	0.03MG; 1.5MG	A201585 001	Nov 18, 2015
	TABLET, ORALLY DISINTEGRATING; ORAL			
	FEMLYV			
	+! MILLICENT PR	0.02MG; 1MG	N218718 001	Jul 22, 2024

ETHINYL ESTRADIOL; NORGESTIMATE

TABLET; ORAL-28

ESTARYLLA

<u>AB</u>	XIROMED	<u>0.035MG; 0.25MG</u>	<u>A090794 001</u>	Jan 30, 2013
	<u>MILI</u>			
<u>AB</u>	AUROBINDO PHARMA LTD	<u>0.035MG; 0.25MG</u>	<u>A205449 001</u>	Jul 07, 2016
	<u>MONO-LINYAH</u>			
<u>AB</u>	NOVAST LABS LTD	<u>0.035MG; 0.25MG</u>	<u>A090523 001</u>	May 23, 2012
	<u>NORGESTIMATE AND ETHINYL ESTRADIOL</u>			
<u>AB</u>	AMNEAL PHARMS	<u>0.025MG, 0.025MG, 0.025MG; 0.18MG, 0.215MG, 0.25MG</u>	<u>A203873 001</u>	May 12, 2016
<u>AB</u>	! GLENMARK PHARMS LTD	<u>0.035MG, 0.035MG, 0.035MG; 0.18MG, 0.215MG, 0.25MG</u>	<u>A200494 001</u>	Jun 17, 2011
<u>AB</u>		<u>0.035MG; 0.25MG</u>	<u>A200538 001</u>	Apr 05, 2012
<u>AB</u>	! GLENMARK SPECLT	<u>0.025MG, 0.025MG, 0.025MG; 0.18MG, 0.215MG, 0.25MG</u>	<u>A204057 001</u>	Feb 23, 2016
<u>AB</u>	LUPIN LTD	<u>0.035MG, 0.035MG, 0.035MG; 0.18MG, 0.215MG, 0.25MG</u>	<u>A205588 001</u>	Apr 26, 2016
<u>AB</u>		<u>0.035MG; 0.25MG</u>	<u>A205630 001</u>	Oct 27, 2016
<u>AB</u>	LUPIN PHARMS	<u>0.025MG, 0.025MG, 0.025MG; 0.18MG, 0.215MG, 0.25MG</u>	<u>A200541 001</u>	Jun 25, 2012
<u>AB</u>	NAARI PTE LTD	<u>0.035MG, 0.035MG, 0.035MG; 0.18MG, 0.215MG, 0.25MG</u>	<u>A200383 001</u>	Apr 07, 2015
<u>AB</u>		<u>0.035MG; 0.25MG</u>	<u>A200384 001</u>	Apr 07, 2015
	<u>PREVIFEM</u>			
<u>AB</u>	ENDO OPERATIONS	<u>0.035MG; 0.25MG</u>	<u>A076334 001</u>	Jan 09, 2004
	<u>SPRINTEC</u>			
<u>AB</u>	! BARR	<u>0.035MG; 0.25MG</u>	<u>A075804 001</u>	Sep 25, 2002
	<u>TRI-LO SPRINTEC</u>			
<u>AB</u>	BARR LABS INC	<u>0.025MG, 0.025MG, 0.025MG; 0.18MG, 0.215MG, 0.25MG</u>	<u>A076784 001</u>	Jun 29, 2009
	<u>TRI-ESTARYLLA</u>			
<u>AB</u>	XIROMED	<u>0.035MG, 0.035MG, 0.035MG; 0.18MG, 0.215MG, 0.25MG</u>	<u>A090793 001</u>	Jan 30, 2013
	<u>TRI-LINYAH</u>			
<u>AB</u>	NOVAST LABS LTD	<u>0.035MG, 0.035MG, 0.035MG; 0.18MG, 0.215MG, 0.25MG</u>	<u>A090524 001</u>	May 30, 2012
	<u>TRI-LO-ESTARYLLA</u>			
<u>AB</u>	XIROMED	<u>0.025MG, 0.025MG, 0.025MG; 0.18MG, 0.215MG, 0.25MG</u>	<u>A091232 001</u>	Jun 29, 2015
	<u>TRI-LO-LINYAH</u>			
<u>AB</u>	NOVAST LABS	<u>0.025MG, 0.025MG, 0.025MG; 0.18MG, 0.215MG, 0.25MG</u>	<u>A090541 001</u>	Sep 02, 2022
	<u>TRI-LO-MILI</u>			
<u>AB</u>	AUROBINDO PHARMA	<u>0.025MG, 0.025MG, 0.025MG; 0.18MG, 0.215MG, 0.25MG</u>	<u>A205762 001</u>	Nov 04, 2016
	<u>TRI-MILI</u>			
<u>AB</u>	AUROBINDO PHARMA LTD	<u>0.035MG, 0.035MG, 0.035MG; 0.18MG, 0.215MG, 0.25MG</u>	<u>A205441 001</u>	Jul 06, 2016
	<u>TRI-SPRINTEC</u>			
<u>AB</u>	BARR	<u>0.035MG, 0.035MG, 0.035MG; 0.18MG, 0.215MG, 0.25MG</u>	<u>A075808 001</u>	Dec 29, 2003

ETHINYL ESTRADIOL; NORGESTREL

TABLET; ORAL-21

CRYSELLE

<u>AB</u>	DURAMED PHARMS BARR	<u>0.03MG; 0.3MG</u>	<u>A075840 001</u>	Nov 30, 2001
	TABLET; ORAL-28			
	<u>CRYSELLE</u>			
<u>AB</u>	! DURAMED PHARMS BARR	<u>0.03MG; 0.3MG</u>	<u>A075840 002</u>	Nov 30, 2001
	<u>ELINEST</u>			
<u>AB</u>	NOVAST LABS LTD	<u>0.03MG; 0.3MG</u>	<u>A091105 001</u>	Mar 28, 2012

PRESCRIPTION DRUG PRODUCT LIST

ETHINYL ESTRADIOL; NORGESTREL

TABLET; ORAL-28

LOW-OGESTREL-28**AB** DR REDDYS LABS SA **0.03MG;0.3MG** **A075288 002** Jul 28, 1999TURQOZ**AB** LUPIN LTD **0.03MG;0.3MG** **A202980 001** Jul 31, 2023ETHINYL ESTRADIOL; SEGESTERONE ACETATE

RING; VAGINAL

ANNOVERA

+! MAYNE PHARMA 0.013MG/24HR;0.15MG/24HR N209627 001 Aug 10, 2018

ETHIODIZED OIL

OIL; INTRALYMPHATIC, INTRAUTERINE

LIPIODOL

+! GUERBET EQ 4.8GM IODINE/10ML (EQ 480MG IODINE/ML) N009190 001

ETHIONAMIDE

TABLET; ORAL

TRECATOR

+! WYETH PHARMS 250MG N013026 002

ETHOSUXIMIDE

CAPSULE; ORAL

ETHOSUXIMIDE**AB** BIONPHARMA **250MG** **A040430 001** Oct 28, 2002**AB** EPIC PHARMA LLC **250MG** **A040686 001** May 28, 2008**AB** HERITAGE PHARMS INC **250MG** **A200892 001** Sep 25, 2012**AB** PURACAP PHARM LLC **250MG** **A210654 001** Mar 16, 2020**AB** STRIDES SOFTGELS **250MG** **A211928 001** Feb 19, 2019ZARONTIN**AB** +! PARKE DAVIS **250MG** **N012380 001**

SYRUP; ORAL

ETHOSUXIMIDE**AA** MIKART **250MG/5ML** **A040506 001** Dec 22, 2003**AA** PHARM ASSOC **250MG/5ML** **A040253 001** Nov 22, 2000ZARONTIN**AA** +! PARKE-DAVIS **250MG/5ML** **A080258 001**ETODOLAC

CAPSULE; ORAL

ETODOLAC**AB** ANI PHARMS **200MG** **A075126 001** Sep 16, 1999**AB** **300MG** **A075126 002** Sep 16, 1999**AB** APOTEX **200MG** **A075419 001** Jul 28, 2000**AB** **300MG** **A075419 002** Jul 28, 2000**AB** IPCA LABS LTD **200MG** **A205448 001** Aug 01, 2024**AB** **300MG** **A205448 002** Aug 01, 2024**AB** TARO **200MG** **A075078 001** Apr 30, 1998**AB** ! **300MG** **A075078 002** Apr 30, 1998

TABLET; ORAL

ETODOLAC**AB** ADAPTIS **400MG** **A209888 001** Nov 30, 2018**AB** **500MG** **A209888 002** Nov 30, 2018**AB** AMNEAL PHARMS CO **400MG** **A208834 001** Jun 07, 2018**AB** **500MG** **A208834 002** Jun 07, 2018**AB** APOTEX INC **400MG** **A076004 001** Dec 03, 2002**AB** **500MG** **A076004 002** Dec 03, 2002**AB** IPCA LABS LTD **400MG** **A204729 001** May 01, 2024**AB** **500MG** **A204729 002** May 01, 2024**AB** PANGEA **400MG** **A074903 001** Apr 11, 1997**AB** **500MG** **A074903 002** Apr 19, 1999**AB** TARO PHARM INDS **400MG** **A075074 001** Mar 11, 1998**AB** ! **500MG** **A075074 002** Apr 25, 2000**AB** UNICHEM **400MG** **A210704 001** Dec 16, 2020**AB** **500MG** **A210704 002** Dec 16, 2020

TABLET, EXTENDED RELEASE; ORAL

ETODOLAC**AB** TARO **400MG** **A076174 001** Mar 13, 2003**AB** **500MG** **A076174 002** Mar 13, 2003**AB** **600MG** **A076174 003** Mar 13, 2003**AB** TEVA **400MG** **A075665 003** Feb 05, 2001**AB** **500MG** **A075665 002** Jul 31, 2000**AB** ! **600MG** **A075665 001** Jul 31, 2000

PRESCRIPTION DRUG PRODUCT LIST

ETODOLAC

TABLET, EXTENDED RELEASE;ORAL

ETODOLAC

<u>AB</u>	UNICHEM	<u>400MG</u>	<u>A212263 001</u>	Nov 24, 2020
<u>AB</u>		<u>500MG</u>	<u>A212263 002</u>	Nov 24, 2020
<u>AB</u>		<u>600MG</u>	<u>A212263 003</u>	Nov 24, 2020
<u>AB</u>	ZYDUS PHARMS	<u>400MG</u>	<u>A091134 001</u>	Jan 23, 2014
<u>AB</u>		<u>500MG</u>	<u>A091134 002</u>	Jan 23, 2014
<u>AB</u>		<u>600MG</u>	<u>A091134 003</u>	Jan 23, 2014

ETOMIDATE

INJECTABLE; INJECTION

AMIDATE

<u>AP</u>	+!	HOSPIRA	<u>2MG/ML</u>	<u>N018227 001</u>	Sep 07, 1982
-----------	----	---------	---------------	--------------------	--------------

ETOMIDATE

<u>AP</u>		CAPLIN	<u>2MG/ML</u>	<u>A215028 001</u>	Dec 18, 2020
<u>AP</u>		EUGIA PHARMA	<u>2MG/ML</u>	<u>A206126 001</u>	Feb 24, 2017
<u>AP</u>		GLAND PHARMA LTD	<u>2MG/ML</u>	<u>A209058 001</u>	Apr 18, 2017
<u>AP</u>		HIKMA	<u>2MG/ML</u>	<u>A074593 001</u>	Nov 04, 1996
<u>AP</u>			<u>2MG/ML</u>	<u>A202354 001</u>	Feb 25, 2016
<u>AP</u>		MYLAN LABS LTD	<u>2MG/ML</u>	<u>A201044 001</u>	Feb 07, 2017
<u>AP</u>		ZYDUS PHARMS	<u>2MG/ML</u>	<u>A202360 001</u>	Jul 18, 2014

ETONOGESTREL

IMPLANT; IMPLANTATION

NEXPLANON

+!	ORGANON	68MG/IMPLANT	N021529 002	May 13, 2011
----	---------	--------------	-------------	--------------

ETOPOSIDE

CAPSULE; ORAL

ETOPOSIDE

!	MYLAN	50MG	A075635 001	Sep 19, 2001
---	-------	------	-------------	--------------

INJECTABLE; INJECTION

ETOPOSIDE

<u>AP</u>		ACCORD HLTHCARE	<u>20MG/ML</u>	<u>A074513 001</u>	Mar 14, 1996
<u>AP</u>	!	FRESENIUS KABI USA	<u>20MG/ML</u>	<u>A074983 001</u>	Sep 30, 1998
<u>AP</u>		HIKMA	<u>20MG/ML</u>	<u>A074290 001</u>	Jul 17, 1995
<u>AP</u>		MEITHEAL	<u>20MG/ML</u>	<u>A074529 001</u>	Jul 24, 1996

ETOPOSIDE PHOSPHATE

INJECTABLE; INJECTION

ETOPHOS PRESERVATIVE FREE

+!	CHEPLAPHARM	EQ 100MG BASE/VIAL	N020457 001	May 17, 1996
----	-------------	--------------------	-------------	--------------

ETRASIMOD ARGININE

TABLET; ORAL

VELSIPITY

+!	PFIZER	EQ 2MG BASE	N216956 001	Oct 12, 2023
----	--------	-------------	-------------	--------------

ETRAVIRINE

TABLET; ORAL

ETRAVIRINE

<u>AB</u>		AMNEAL	<u>100MG</u>	<u>A214196 002</u>	Jun 14, 2021
<u>AB</u>			<u>200MG</u>	<u>A214196 003</u>	Jun 14, 2021
<u>AB</u>		CARNEGIE	<u>100MG</u>	<u>A215402 001</u>	Apr 13, 2022
<u>AB</u>			<u>200MG</u>	<u>A215402 002</u>	Apr 13, 2022

INTELENCE

<u>AB</u>	+	JANSSEN R AND D	<u>100MG</u>	<u>N022187 001</u>	Jan 18, 2008
<u>AB</u>	+!		<u>200MG</u>	<u>N022187 002</u>	Dec 22, 2010
	+		25MG	N022187 003	Mar 26, 2012

EVEROLIMUS

TABLET; ORAL

AFINITOR

<u>AB</u>	+	NOVARTIS	<u>2.5MG</u>	<u>N022334 003</u>	Jul 09, 2010
<u>AB</u>	+!		<u>5MG</u>	<u>N022334 001</u>	Mar 30, 2009
<u>AB</u>	+		<u>7.5MG</u>	<u>N022334 004</u>	Mar 30, 2012
<u>AB</u>	+		<u>10MG</u>	<u>N022334 002</u>	Mar 30, 2009

EVEROLIMUS

<u>AB</u>		ALKEM LABS LTD	<u>0.25MG</u>	<u>A214138 001</u>	Nov 26, 2021
<u>AB</u>			<u>0.5MG</u>	<u>A214138 002</u>	Nov 26, 2021
<u>AB</u>			<u>0.75MG</u>	<u>A214138 003</u>	Nov 26, 2021
<u>AB</u>			<u>1MG</u>	<u>A214138 004</u>	Nov 26, 2021
<u>AB</u>		BIOCON PHARMA	<u>2.5MG</u>	<u>A214182 001</u>	Feb 11, 2021
<u>AB</u>			<u>5MG</u>	<u>A214182 002</u>	Feb 11, 2021
<u>AB</u>			<u>7.5MG</u>	<u>A214182 003</u>	Feb 11, 2021

PRESCRIPTION DRUG PRODUCT LIST

EVEROLIMUS

TABLET; ORAL

EVEROLIMUS

<u>AB</u>		<u>10MG</u>	<u>A214182</u>	<u>004</u>	Feb 11, 2021
<u>AB</u>	BRECKENRIDGE	<u>0.25MG</u>	<u>A205432</u>	<u>001</u>	May 20, 2021
<u>AB</u>		<u>0.5MG</u>	<u>A205432</u>	<u>002</u>	May 20, 2021
<u>AB</u>		<u>0.75MG</u>	<u>A205432</u>	<u>003</u>	May 20, 2021
<u>AB</u>		<u>2.5MG</u>	<u>A205426</u>	<u>001</u>	Mar 05, 2021
<u>AB</u>		<u>5MG</u>	<u>A205426</u>	<u>002</u>	Mar 05, 2021
<u>AB</u>		<u>7.5MG</u>	<u>A205426</u>	<u>003</u>	Mar 05, 2021
<u>AB</u>		<u>10MG</u>	<u>A205426</u>	<u>004</u>	Mar 05, 2021
<u>AB</u>	ENDO OPERATIONS	<u>0.25MG</u>	<u>A205775</u>	<u>001</u>	Oct 18, 2021
<u>AB</u>		<u>0.5MG</u>	<u>A205775</u>	<u>002</u>	Oct 18, 2021
<u>AB</u>		<u>0.75MG</u>	<u>A205775</u>	<u>003</u>	Oct 18, 2021
<u>AB</u>		<u>1MG</u>	<u>A205775</u>	<u>004</u>	Oct 18, 2021
<u>AB</u>		<u>2.5MG</u>	<u>A207934</u>	<u>001</u>	Dec 09, 2019
<u>AB</u>		<u>5MG</u>	<u>A207934</u>	<u>002</u>	Dec 09, 2019
<u>AB</u>		<u>7.5MG</u>	<u>A207934</u>	<u>003</u>	Dec 09, 2019
<u>AB</u>		<u>10MG</u>	<u>A207934</u>	<u>004</u>	Dec 09, 2020
<u>AB</u>	HIKMA	<u>0.25MG</u>	<u>A206133</u>	<u>001</u>	Apr 12, 2018
<u>AB</u>		<u>0.5MG</u>	<u>A206133</u>	<u>002</u>	Apr 12, 2018
<u>AB</u>		<u>0.75MG</u>	<u>A206133</u>	<u>003</u>	Apr 12, 2018
<u>AB</u>		<u>1MG</u>	<u>A206133</u>	<u>004</u>	Nov 18, 2021
<u>AB</u>		<u>2.5MG</u>	<u>A207486</u>	<u>001</u>	Jun 08, 2020
<u>AB</u>		<u>5MG</u>	<u>A207486</u>	<u>002</u>	Jun 08, 2020
<u>AB</u>		<u>7.5MG</u>	<u>A207486</u>	<u>003</u>	Jun 08, 2020
<u>AB</u>		<u>10MG</u>	<u>A207486</u>	<u>004</u>	Nov 23, 2021
<u>AB</u>	MYLAN	<u>2.5MG</u>	<u>A212936</u>	<u>001</u>	Jun 08, 2020
<u>AB</u>		<u>5MG</u>	<u>A212936</u>	<u>002</u>	Jun 08, 2020
<u>AB</u>		<u>7.5MG</u>	<u>A212936</u>	<u>003</u>	Jun 08, 2020
<u>AB</u>		<u>10MG</u>	<u>A212936</u>	<u>004</u>	Jun 08, 2020
<u>AB</u>	TEVA PHARMS USA	<u>2.5MG</u>	<u>A210050</u>	<u>001</u>	Dec 09, 2019
<u>AB</u>		<u>5MG</u>	<u>A210050</u>	<u>002</u>	Dec 09, 2019
<u>AB</u>		<u>7.5MG</u>	<u>A210050</u>	<u>003</u>	Dec 09, 2019
<u>AB</u>		<u>10MG</u>	<u>A210050</u>	<u>004</u>	Dec 09, 2019

ZORTRESS

<u>AB</u>	+	NOVARTIS	<u>0.25MG</u>	<u>N021560</u>	<u>001</u>	Apr 20, 2010
<u>AB</u>	+		<u>0.5MG</u>	<u>N021560</u>	<u>002</u>	Apr 20, 2010
<u>AB</u>	+		<u>0.75MG</u>	<u>N021560</u>	<u>003</u>	Apr 20, 2010
<u>AB</u>	+	!	<u>1MG</u>	<u>N021560</u>	<u>004</u>	Aug 10, 2018

TABLET, FOR SUSPENSION; ORAL

AFINITOR DISPERZ

<u>AB</u>	+	NOVARTIS PHARM	<u>2MG</u>	<u>N203985</u>	<u>001</u>	Aug 29, 2012
<u>AB</u>	+		<u>3MG</u>	<u>N203985</u>	<u>002</u>	Aug 29, 2012
<u>AB</u>	+	!	<u>5MG</u>	<u>N203985</u>	<u>003</u>	Aug 29, 2012

EVEROLIMUS

<u>AB</u>		MYLAN	<u>2MG</u>	<u>A210130</u>	<u>001</u>	Apr 19, 2019
<u>AB</u>			<u>3MG</u>	<u>A210130</u>	<u>002</u>	Apr 19, 2019
<u>AB</u>			<u>5MG</u>	<u>A210130</u>	<u>003</u>	Apr 19, 2019

EXEMESTANE

TABLET; ORAL

AROMASIN

<u>AB</u>	+	!	PFIZER	<u>25MG</u>	<u>N020753</u>	<u>001</u>	Oct 21, 1999
-----------	---	---	--------	-------------	----------------	------------	--------------

EXEMESTANE

<u>AB</u>		BRECKENRIDGE	<u>25MG</u>	<u>A211031</u>	<u>001</u>	Feb 21, 2019
<u>AB</u>		CIPLA	<u>25MG</u>	<u>A210323</u>	<u>001</u>	Apr 27, 2018
<u>AB</u>		EUGIA PHARMA	<u>25MG</u>	<u>A216454</u>	<u>001</u>	May 20, 2022
<u>AB</u>		HIKMA	<u>25MG</u>	<u>A077431</u>	<u>001</u>	Apr 01, 2011
<u>AB</u>		QILU	<u>25MG</u>	<u>A213547</u>	<u>001</u>	Apr 13, 2020
<u>AB</u>		RISING	<u>25MG</u>	<u>A203315</u>	<u>001</u>	Mar 10, 2017
<u>AB</u>		UPSHER SMITH LABS	<u>25MG</u>	<u>A209208</u>	<u>001</u>	Jul 26, 2017
<u>AB</u>		ZYDUS PHARMS	<u>25MG</u>	<u>A202602</u>	<u>001</u>	Oct 03, 2018

EXENATIDE SYNTHETIC

INJECTABLE; SUBCUTANEOUS

BYETTA

<u>AP</u>	+	!	ASTRAZENECA AB	<u>300MCG/1.2ML (250MCG/ML)</u>	<u>N021773</u>	<u>001</u>	Apr 28, 2005
<u>AP</u>	+	!		<u>600MCG/2.4ML (250MCG/ML)</u>	<u>N021773</u>	<u>002</u>	Apr 28, 2005

EXENATIDE SYNTHETIC

<u>AP</u>		AMNEAL	<u>300MCG/1.2ML (250MCG/ML)</u>	<u>A206697</u>	<u>001</u>	Nov 19, 2024
<u>AP</u>			<u>600MCG/2.4ML (250MCG/ML)</u>	<u>A206697</u>	<u>002</u>	Nov 19, 2024

PRESCRIPTION DRUG PRODUCT LISTEXENATIDE SYNTHETIC

SUSPENSION, EXTENDED RELEASE;SUBCUTANEOUS

BYDUREON BCISE

+! ASTRAZENECA AB 2MG/0.85ML (2MG/0.85ML) N209210 001 Oct 20, 2017

EZETIMIBE

TABLET;ORAL

EZETIMIBE

<u>AB</u>	ACCORD HLTHCARE	<u>10MG</u>	<u>A211550</u>	<u>001</u>	Oct 26, 2018
<u>AB</u>	ALKEM LABS LTD	<u>10MG</u>	<u>A209234</u>	<u>001</u>	Dec 21, 2017
<u>AB</u>	AMNEAL PHARMS CO	<u>10MG</u>	<u>A208803</u>	<u>001</u>	Jun 12, 2017
<u>AB</u>	AUROBINDO PHARMA	<u>10MG</u>	<u>A209838</u>	<u>001</u>	Aug 25, 2017
<u>AB</u>	GLENMARK PHARMS LTD	<u>10MG</u>	<u>A078560</u>	<u>001</u>	Jun 26, 2015
<u>AB</u>	HETERO LABS LTD III	<u>10MG</u>	<u>A210859</u>	<u>001</u>	Jul 26, 2022
<u>AB</u>	MACLEODS PHARMS LTD	<u>10MG</u>	<u>A211159</u>	<u>001</u>	Apr 29, 2024
<u>AB</u>	OHM LABS INC	<u>10MG</u>	<u>A207311</u>	<u>001</u>	Jun 12, 2017
<u>AB</u>	ORIENT PHARMA	<u>10MG</u>	<u>A215693</u>	<u>001</u>	Sep 13, 2022
<u>AB</u>	SANDOZ	<u>10MG</u>	<u>A203931</u>	<u>001</u>	Jun 12, 2017
<u>AB</u>	SCIEGEN PHARMS INC	<u>10MG</u>	<u>A210673</u>	<u>001</u>	Oct 23, 2020
<u>AB</u>	WATSON LABS INC	<u>10MG</u>	<u>A200831</u>	<u>001</u>	Jun 12, 2017
<u>AB</u>	ZYDUS PHARMS	<u>10MG</u>	<u>A204331</u>	<u>001</u>	Jun 12, 2017

ZETIA

<u>AB</u>	+! ORGANON	<u>10MG</u>	<u>N021445</u>	<u>001</u>	Oct 25, 2002
-----------	------------	-------------	----------------	------------	--------------

EZETIMIBE; SIMVASTATIN

TABLET;ORAL

EZETIMIBE AND SIMVASTATIN

<u>AB</u>	ALKEM LABS LTD	<u>10MG;10MG</u>	<u>A209222</u>	<u>001</u>	Dec 22, 2017
<u>AB</u>		<u>10MG;20MG</u>	<u>A209222</u>	<u>002</u>	Dec 22, 2017
<u>AB</u>		<u>10MG;40MG</u>	<u>A209222</u>	<u>003</u>	Dec 22, 2017
<u>AB</u>		<u>10MG;80MG</u>	<u>A209222</u>	<u>004</u>	Dec 22, 2017
<u>AB</u>	AMNEAL PHARMS CO	<u>10MG;10MG</u>	<u>A208831</u>	<u>001</u>	Nov 21, 2017
<u>AB</u>		<u>10MG;20MG</u>	<u>A208831</u>	<u>002</u>	Nov 21, 2017
<u>AB</u>		<u>10MG;40MG</u>	<u>A208831</u>	<u>003</u>	Nov 21, 2017
<u>AB</u>		<u>10MG;80MG</u>	<u>A208831</u>	<u>004</u>	Nov 21, 2017
<u>AB</u>	AUROBINDO PHARMA USA	<u>10MG;10MG</u>	<u>A200082</u>	<u>001</u>	Dec 17, 2020
<u>AB</u>		<u>10MG;20MG</u>	<u>A200082</u>	<u>002</u>	Dec 17, 2020
<u>AB</u>		<u>10MG;40MG</u>	<u>A200082</u>	<u>003</u>	Dec 17, 2020
<u>AB</u>		<u>10MG;80MG</u>	<u>A200082</u>	<u>004</u>	Dec 17, 2020
<u>AB</u>	DR REDDYS LABS SA	<u>10MG;10MG</u>	<u>A200909</u>	<u>001</u>	Apr 26, 2017
<u>AB</u>		<u>10MG;20MG</u>	<u>A200909</u>	<u>002</u>	Apr 26, 2017
<u>AB</u>		<u>10MG;40MG</u>	<u>A200909</u>	<u>003</u>	Apr 26, 2017
<u>AB</u>		<u>10MG;80MG</u>	<u>A200909</u>	<u>004</u>	Apr 26, 2017
<u>AB</u>	GLENMARK PHARMS LTD	<u>10MG;10MG</u>	<u>A208699</u>	<u>001</u>	Jun 27, 2019
<u>AB</u>		<u>10MG;20MG</u>	<u>A208699</u>	<u>002</u>	Jun 27, 2019
<u>AB</u>		<u>10MG;40MG</u>	<u>A208699</u>	<u>003</u>	Jun 27, 2019
<u>AB</u>		<u>10MG;80MG</u>	<u>A208699</u>	<u>004</u>	Jun 27, 2019
<u>AB</u>	SCIEGEN PHARMS INC	<u>10MG;10MG</u>	<u>A211663</u>	<u>001</u>	Dec 10, 2024
<u>AB</u>		<u>10MG;20MG</u>	<u>A211663</u>	<u>002</u>	Dec 10, 2024
<u>AB</u>		<u>10MG;40MG</u>	<u>A211663</u>	<u>003</u>	Dec 10, 2024
<u>AB</u>		<u>10MG;80MG</u>	<u>A211663</u>	<u>004</u>	Dec 10, 2024
<u>AB</u>	WATSON LABS INC	<u>10MG;10MG</u>	<u>A202968</u>	<u>001</u>	Apr 26, 2017
<u>AB</u>		<u>10MG;20MG</u>	<u>A202968</u>	<u>002</u>	Apr 26, 2017
<u>AB</u>		<u>10MG;40MG</u>	<u>A202968</u>	<u>003</u>	Apr 26, 2017
<u>AB</u>		<u>10MG;80MG</u>	<u>A202968</u>	<u>004</u>	Apr 26, 2017

VYTORIN

<u>AB</u>	+ ORGANON	<u>10MG;10MG</u>	<u>N021687</u>	<u>001</u>	Jul 23, 2004
<u>AB</u>	+	<u>10MG;20MG</u>	<u>N021687</u>	<u>002</u>	Jul 23, 2004
<u>AB</u>	+	<u>10MG;40MG</u>	<u>N021687</u>	<u>003</u>	Jul 23, 2004
<u>AB</u>	+!	<u>10MG;80MG</u>	<u>N021687</u>	<u>004</u>	Jul 23, 2004

FAMCICLOVIR

TABLET;ORAL

FAMCICLOVIR

<u>AB</u>	APOTEX	<u>125MG</u>	<u>A091480</u>	<u>001</u>	Jul 22, 2011
<u>AB</u>		<u>250MG</u>	<u>A091480</u>	<u>002</u>	Jul 22, 2011
<u>AB</u>		<u>500MG</u>	<u>A091480</u>	<u>003</u>	Jul 22, 2011
<u>AB</u>	AUROBINDO PHARMA LTD	<u>125MG</u>	<u>A091114</u>	<u>001</u>	Mar 21, 2011
<u>AB</u>		<u>250MG</u>	<u>A091114</u>	<u>002</u>	Mar 21, 2011
<u>AB</u>		<u>500MG</u>	<u>A091114</u>	<u>003</u>	Mar 21, 2011
<u>AB</u>	HETERO LABS LTD V	<u>125MG</u>	<u>A202438</u>	<u>001</u>	Sep 10, 2014
<u>AB</u>		<u>250MG</u>	<u>A202438</u>	<u>002</u>	Sep 10, 2014

PRESCRIPTION DRUG PRODUCT LIST

FAMCICLOVIR

TABLET; ORAL

FAMCICLOVIR

AB		500MG	A202438 003	Sep 10, 2014
AB	MACLEODS PHARMS LTD	125MG	A201022 001	Jan 12, 2012
AB		250MG	A201022 002	Jan 12, 2012
AB		500MG	A201022 003	Jan 12, 2012
AB	TEVA PHARMS	125MG	A077487 001	Aug 24, 2007
AB		250MG	A077487 002	Aug 24, 2007
AB	!	500MG	A077487 003	Aug 24, 2007

FAMOTIDINE

FOR SUSPENSION; ORAL

FAMOTIDINE

AB	AJANTA PHARMA LTD	40MG/5ML	A217529 001	Sep 18, 2023
AB	AKORN	40MG/5ML	A201995 001	May 30, 2014
AB	ALKEM LABS LTD	40MG/5ML	A216400 001	May 15, 2023
AB	AMNEALS PHARMS	40MG/5ML	A216427 001	Aug 04, 2022
AB	ANNORA PHARMA	40MG/5ML	A217330 001	Aug 17, 2023
AB	CARNEGIE	40MG/5ML	A217137 001	Jul 07, 2023
AB	EPIC PHARMA LLC	40MG/5ML	A218344 001	Oct 16, 2024
AB	LUPIN LTD	40MG/5ML	A090440 001	Jun 29, 2010
AB	MICRO LABS	40MG/5ML	A217842 001	Sep 14, 2023
AB	NAVINTA LLC	40MG/5ML	A091020 001	May 27, 2010
AB	NOVEL LABS INC	40MG/5ML	A201695 001	Dec 17, 2012
AB	! NOVIITIUM PHARMA	40MG/5ML	A215043 001	Apr 20, 2021
AB	UNICHEM	40MG/5ML	A217605 001	Jan 16, 2024
AB	UPSHER SMITH LABS	40MG/5ML	A217655 001	Jun 16, 2023

INJECTABLE; INJECTION

FAMOTIDINE

AP	FRESENIUS KABI USA	10MG/ML	A075709 001	Apr 16, 2001
AP	! HIKMA	10MG/ML	A075488 001	Apr 16, 2001
AP	MYLAN LABS LTD	10MG/ML	A078641 001	Jun 25, 2008
AP	SAGENT	10MG/ML	A075651 001	Apr 16, 2001
AP		10MG/ML	A075684 001	Apr 16, 2001

FAMOTIDINE PRESERVATIVE FREE

AP	FRESENIUS KABI USA	10MG/ML	A075813 001	Apr 16, 2001
AP	! HIKMA	10MG/ML	A075486 001	Apr 16, 2001
AP	MYLAN LABS LTD	10MG/ML	A078642 001	Jun 25, 2008
AP	SAGENT	10MG/ML	A075622 001	Apr 16, 2001
AP		10MG/ML	A075825 001	Apr 17, 2001

FAMOTIDINE PRESERVATIVE FREE IN PLASTIC CONTAINER

!	BAXTER HLTHCARE	0.4MG/ML	A075591 001	May 10, 2001
---	-----------------	----------	-------------	--------------

TABLET; ORAL

FAMOTIDINE

AB	ALEMBIC PHARMS LTD	20MG	A078916 001	May 22, 2009
AB		40MG	A078916 002	May 22, 2009
AB	ALKEM LABS LTD	20MG	A215630 001	Jan 07, 2022
AB		20MG	A217375 001	Apr 24, 2023
AB		40MG	A215630 002	Jan 07, 2022
AB		40MG	A217375 002	Apr 24, 2023
AB	ANNORA PHARMA	20MG	A215767 001	Nov 04, 2021
AB		40MG	A215767 002	Nov 04, 2021
AB	APOTEX	20MG	A075611 001	Jul 23, 2001
AB		40MG	A075611 002	Jul 23, 2001
AB	ASCENT PHARMS INC	20MG	A215689 001	Oct 15, 2021
AB		40MG	A215689 002	Oct 15, 2021
AB	AUROBINDO PHARMA LTD	20MG	A206530 001	Dec 22, 2015
AB	!	40MG	A206530 002	Dec 22, 2015
AB	CARLSBAD	20MG	A075805 001	Apr 16, 2001
AB		40MG	A075805 002	Apr 16, 2001
AB	CHARTWELL RX	20MG	A075786 001	Apr 16, 2001
AB		40MG	A075786 002	Apr 16, 2001
AB	CONTRACT PHARMACAL	20MG	A217669 001	Dec 20, 2023
AB		40MG	A217669 002	Dec 20, 2023
AB	DR REDDYS LABS LTD	20MG	A075718 001	Apr 16, 2001
AB		40MG	A075718 002	Apr 16, 2001
AB	IVAX SUB TEVA PHARMS	20MG	A075511 001	Apr 16, 2001
AB		40MG	A075511 002	Apr 16, 2001
AB	MANKIND PHARMA	20MG	A075302 001	Apr 16, 2001
AB		40MG	A075302 002	Apr 16, 2001

PRESCRIPTION DRUG PRODUCT LIST

FAMOTIDINE

TABLET; ORAL

FAMOTIDINE

<u>AB</u>	MSN	<u>20MG</u>	<u>A217400 001</u>	Aug 06, 2024
<u>AB</u>		<u>40MG</u>	<u>A217400 002</u>	Aug 06, 2024
<u>AB</u>	RISING	<u>20MG</u>	<u>A218181 001</u>	Dec 22, 2023
<u>AB</u>		<u>40MG</u>	<u>A218181 002</u>	Dec 22, 2023
<u>AB</u>	ZYDUS PHARMS	<u>20MG</u>	<u>A216441 001</u>	Jun 03, 2022
<u>AB</u>		<u>40MG</u>	<u>A216441 002</u>	Jun 03, 2022

FAMOTIDINE; IBUPROFEN

TABLET; ORAL

FAMOTIDINE; IBUPROFEN

<u>AB</u>	ENDO OPERATIONS	<u>26.6MG;800MG</u>	<u>A203658 001</u>	Mar 22, 2024
-----------	-----------------	---------------------	--------------------	--------------

IBUPROFEN AND FAMOTIDINE

<u>AB</u>	! ALKEM LABS LTD	<u>26.6MG;800MG</u>	<u>A211890 001</u>	Aug 03, 2021
<u>AB</u>	ASCENT PHARMS INC	<u>26.6MG;800MG</u>	<u>A216814 001</u>	Mar 15, 2023
<u>AB</u>	TEVA PHARMS USA	<u>26.6MG;800MG</u>	<u>A211278 001</u>	Oct 29, 2021

FEBUXOSTAT

TABLET; ORAL

FEBUXOSTAT

<u>AB</u>	ALEMBIC	<u>40MG</u>	<u>A205421 001</u>	Jul 01, 2019
<u>AB</u>		<u>80MG</u>	<u>A205421 002</u>	Jul 01, 2019
<u>AB</u>	ALKEM LABS LTD	<u>40MG</u>	<u>A212924 001</u>	Dec 07, 2021
<u>AB</u>		<u>80MG</u>	<u>A212924 002</u>	Dec 07, 2021
<u>AB</u>	AUROBINDO PHARMA LTD	<u>40MG</u>	<u>A210741 001</u>	Oct 25, 2023
<u>AB</u>		<u>80MG</u>	<u>A210741 002</u>	Oct 25, 2023
<u>AB</u>	DR REDDYS	<u>40MG</u>	<u>A205374 001</u>	Oct 22, 2020
<u>AB</u>		<u>80MG</u>	<u>A205374 002</u>	Oct 22, 2020
<u>AB</u>	HIKMA	<u>40MG</u>	<u>A205414 001</u>	Oct 15, 2019
<u>AB</u>		<u>80MG</u>	<u>A205414 002</u>	Oct 15, 2019
<u>AB</u>	INDOCO	<u>40MG</u>	<u>A210292 001</u>	Dec 30, 2019
<u>AB</u>		<u>80MG</u>	<u>A210292 002</u>	Dec 30, 2019
<u>AB</u>	MACLEODS PHARMS LTD	<u>40MG</u>	<u>A207293 001</u>	Sep 28, 2023
<u>AB</u>		<u>80MG</u>	<u>A207293 002</u>	Sep 28, 2023
<u>AB</u>	PRINSTON INC	<u>40MG</u>	<u>A206266 001</u>	Mar 28, 2022
<u>AB</u>		<u>80MG</u>	<u>A206266 002</u>	Mar 28, 2022
<u>AB</u>	SUN PHARM	<u>40MG</u>	<u>A205467 001</u>	Jul 01, 2019
<u>AB</u>		<u>80MG</u>	<u>A205467 002</u>	Jul 01, 2019
<u>AB</u>	SUNSHINE	<u>40MG</u>	<u>A213069 001</u>	Jun 02, 2020
<u>AB</u>		<u>80MG</u>	<u>A213069 002</u>	Jun 02, 2020
<u>AB</u>	ZYDUS LIFESCIENCES	<u>40MG</u>	<u>A205443 001</u>	Jan 09, 2023
<u>AB</u>		<u>80MG</u>	<u>A205443 002</u>	Jan 09, 2023

ULORIC

<u>AB</u>	+ TAKEDA PHARMS USA	<u>40MG</u>	<u>N021856 001</u>	Feb 13, 2009
<u>AB</u>	+!	<u>80MG</u>	<u>N021856 002</u>	Feb 13, 2009

FEBUXOSTAT

BX	MSN	40MG	A210461 001	Dec 30, 2019
BX		80MG	A210461 002	Dec 30, 2019

FEDRATINIB HYDROCHLORIDE

CAPSULE; ORAL

INREBIC

	+! BRISTOL-MYERS	EQ 100MG BASE	N212327 001	Aug 16, 2019
--	------------------	---------------	-------------	--------------

FELBAMATE

SUSPENSION; ORAL

FELBAMATE

<u>AB</u>	AMNEAL PHARMS	<u>600MG/5ML</u>	<u>A202385 001</u>	Dec 16, 2011
<u>AB</u>	NOVITIUM PHARMA	<u>600MG/5ML</u>	<u>A211333 001</u>	May 31, 2019
<u>AB</u>	TARO	<u>600MG/5ML</u>	<u>A206314 001</u>	Jun 16, 2017

FELBATOL

<u>AB</u>	+! MYLAN SPECIALITY LP	<u>600MG/5ML</u>	<u>N020189 003</u>	Jul 29, 1993
-----------	------------------------	------------------	--------------------	--------------

TABLET; ORAL

FELBAMATE

<u>AB</u>	ALVOGEN	<u>400MG</u>	<u>A204595 001</u>	Jan 11, 2016
<u>AB</u>		<u>600MG</u>	<u>A204595 002</u>	Jan 11, 2016
<u>AB</u>	AMNEAL PHARMS	<u>400MG</u>	<u>A201680 001</u>	Sep 13, 2011
<u>AB</u>		<u>600MG</u>	<u>A201680 002</u>	Sep 13, 2011
<u>AB</u>	ANI PHARMS	<u>400MG</u>	<u>A202284 001</u>	Nov 04, 2015
<u>AB</u>		<u>600MG</u>	<u>A202284 002</u>	Nov 04, 2015
<u>AB</u>	TARO	<u>400MG</u>	<u>A207093 001</u>	Apr 20, 2017
<u>AB</u>		<u>600MG</u>	<u>A207093 002</u>	Apr 20, 2017

PRESCRIPTION DRUG PRODUCT LIST

FELBAMATE

TABLET;ORAL

FELBAMATE

AB	ZYDUS LIFESCIENCES	400MG	A208970 001	May 30, 2017
AB		600MG	A208970 002	May 30, 2017

FELBATOL

AB	+ MYLAN SPECIALITY LP	400MG	N020189 001	Jul 29, 1993
AB	+!	600MG	N020189 002	Jul 29, 1993

FELODIPINE

TABLET, EXTENDED RELEASE;ORAL

FELODIPINE

AB	AUROBINDO PHARMA LTD	2.5MG	A203417 001	Jan 17, 2013
AB		5MG	A203417 002	Jan 17, 2013
AB		10MG	A203417 003	Jan 17, 2013
AB	ENDO OPERATIONS	2.5MG	A200815 001	Oct 28, 2011
AB		5MG	A200815 002	Oct 28, 2011
AB		10MG	A200815 003	Oct 28, 2011
AB	GLENMARK PHARMS LTD	2.5MG	A090365 001	Dec 17, 2010
AB		5MG	A090365 002	Dec 17, 2010
AB		10MG	A090365 003	Dec 17, 2010
AB	HERITAGE	2.5MG	A201964 001	Nov 08, 2013
AB		5MG	A201964 002	Nov 08, 2013
AB		10MG	A201964 003	Nov 08, 2013
AB	ORBION PHARMS	2.5MG	A203032 001	May 21, 2015
AB		5MG	A203032 002	May 21, 2015
AB		10MG	A203032 003	May 21, 2015
AB	SUN PHARM INDS LTD	2.5MG	A091200 001	Dec 13, 2013
AB		5MG	A091200 002	Dec 13, 2013
AB		10MG	A091200 003	Dec 13, 2013
AB	TORRENT PHARMS LTD	2.5MG	A202170 001	Nov 28, 2011
AB		5MG	A202170 002	Nov 28, 2011
AB	!	10MG	A202170 003	Nov 28, 2011
AB	YILING	2.5MG	A210847 001	Oct 26, 2018
AB		5MG	A210847 002	Oct 26, 2018
AB		10MG	A210847 003	Oct 26, 2018
AB	YUNG SHIN PHARM	2.5MG	A204800 001	Apr 29, 2019
AB		5MG	A204800 002	Apr 29, 2019
AB		10MG	A204800 003	Apr 29, 2019

FENFLURAMINE HYDROCHLORIDE

SOLUTION;ORAL

FINTEPLA

+!	UCB INC	EQ 2.2MG BASE/ML	N212102 001	Jun 25, 2020
----	---------	------------------	--------------------	--------------

FENOFIBRATE

CAPSULE;ORAL

ANTARA (MICRONIZED)

AB	+ LUPIN	43MG	N021695 001	Nov 30, 2004
AB	+!	130MG	N021695 003	Nov 30, 2004

FENOFIBRATE

AB	SUN PHARM INDS LTD	43MG	A201748 001	Oct 31, 2014
AB		130MG	A201748 002	Oct 31, 2014

FENOFIBRATE (MICRONIZED)

AB	AJANTA PHARMA LTD	67MG	A210705 001	Sep 10, 2018
AB		134MG	A210705 002	Sep 10, 2018
AB	!	200MG	A210705 003	Sep 10, 2018
AB	ALEMBIC	67MG	A213842 001	Oct 19, 2020
AB		134MG	A213842 002	Oct 19, 2020
AB		200MG	A213842 003	Oct 19, 2020
AB	ANI PHARMS	67MG	A209504 001	Apr 30, 2018
AB		134MG	A209504 002	Apr 30, 2018
AB		200MG	A209504 003	Apr 30, 2018
AB	APOTEX	43MG	A202252 001	Jul 26, 2013
AB		130MG	A202252 002	Jul 26, 2013
AB	AUROBINDO PHARMA LTD	67MG	A212232 001	Sep 20, 2021
AB		134MG	A212232 002	Sep 20, 2021
AB		200MG	A212232 003	Sep 20, 2021
AB	CHARTWELL	67MG	A211407 001	Jan 31, 2024
AB		134MG	A211407 002	Jan 31, 2024
AB		200MG	A211407 003	Jan 31, 2024
AB	DR REDDYS LABS SA	43MG	A090859 001	Mar 01, 2012

PRESCRIPTION DRUG PRODUCT LIST

FENOFIBRATE

CAPSULE; ORAL

FENOFIBRATE (MICRONIZED)

AB		130MG	A090859 002	Mar 01, 2012
AB	GLENMARK PHARMS LTD	67MG	A205566 001	Apr 07, 2017
AB		134MG	A205566 002	Apr 07, 2017
AB		200MG	A205566 003	Apr 07, 2017
AB	REYOUNG	67MG	A207805 001	Nov 16, 2017
AB		134MG	A207805 002	Nov 16, 2017
AB		200MG	A207805 003	Nov 16, 2017
AB	RHODES PHARMS	67MG	A075753 001	Sep 03, 2002
AB		134MG	A075753 002	Apr 09, 2002
AB		200MG	A075753 003	Apr 09, 2002
AB	TORRENT	67MG	A210782 001	Jun 26, 2018
AB		134MG	A210782 002	Jun 26, 2018
AB		200MG	A210782 003	Jun 26, 2018

LIPOFEN

+ CIPHER PHARMS INC
+!50MG
150MGN021612 001 Jan 11, 2006
N021612 003 Jan 11, 2006

TABLET; ORAL

FENOFIBRATE

AB	AJANTA PHARMA LTD	54MG	A210138 001	Jul 23, 2018
AB		160MG	A210138 002	Jul 23, 2018
AB	ALEMBIC	48MG	A210476 001	Aug 09, 2019
AB		145MG	A210476 002	Aug 09, 2019
AB	AMNEAL	48MG	A209951 001	Feb 09, 2018
AB		54MG	A209950 001	Mar 19, 2018
AB		145MG	A209951 002	Feb 09, 2018
AB		160MG	A209950 002	Mar 19, 2018
AB	AUROBINDO PHARMA	48MG	A205118 001	May 05, 2016
AB		54MG	A216798 001	Sep 27, 2022
AB	!	145MG	A205118 002	May 05, 2016
AB		160MG	A216798 002	Sep 27, 2022
AB	AUSTARPHARMA	48MG	A208476 001	Feb 10, 2021
AB		54MG	A207803 001	Dec 19, 2017
AB		145MG	A208476 002	Feb 10, 2021
AB		160MG	A207803 002	Dec 19, 2017
AB	CHARTWELL RX	54MG	A209660 001	Feb 11, 2019
AB		160MG	A209660 002	Feb 11, 2019
AB	CIPLA	48MG	A208709 001	Dec 15, 2016
AB		145MG	A208709 002	Dec 15, 2016
AB	DR REDDYS	54MG	A210670 001	Sep 06, 2019
AB		160MG	A210670 002	Sep 06, 2019
AB	HETERO LABS LTD III	48MG	A204598 001	Jul 12, 2016
AB		145MG	A204598 002	Jul 12, 2016
AB	IMPAX LABS	54MG	A076509 001	Mar 26, 2008
AB	!	160MG	A076509 002	Mar 26, 2008
AB	LUPIN LTD	48MG	A090856 001	Dec 23, 2011
AB		54MG	A204019 001	Aug 17, 2015
AB		145MG	A090856 002	Dec 23, 2011
AB		160MG	A204019 002	Aug 17, 2015
AB	MACLEODS PHARMS LTD	48MG	A210248 001	Nov 13, 2024
AB		145MG	A210248 002	Nov 13, 2024
AB	MANKIND PHARMA	54MG	A213864 001	Jun 12, 2020
AB		160MG	A213864 002	Jun 12, 2020
AB	MYLAN	54MG	A076520 001	Oct 25, 2007
AB		160MG	A076520 003	Oct 25, 2007
AB	MYLAN PHARMS INC	40MG	A204475 001	Jun 23, 2016
AB		48MG	A202856 001	Dec 07, 2012
AB	!	120MG	A204475 002	Jun 23, 2016
AB		145MG	A202856 002	Dec 07, 2012
AB	PRINSTON INC	48MG	A211080 001	Aug 28, 2018
AB		145MG	A211080 002	Aug 28, 2018
AB	RHODES PHARMS	54MG	A076433 001	May 13, 2005
AB		160MG	A076433 002	May 13, 2005
AB	RISING	48MG	A211122 001	Mar 18, 2020
AB		54MG	A210606 001	Aug 17, 2018
AB		145MG	A211122 002	Mar 18, 2020
AB		160MG	A210606 002	Aug 17, 2018
AB	SUN PHARM	48MG	A200884 001	Sep 07, 2017
AB		145MG	A200884 002	Sep 07, 2017
AB	VALEANT PHARMS	48MG	A090715 001	Apr 05, 2012

PRESCRIPTION DRUG PRODUCT LIST

FENOFIBRATE

TABLET; ORAL

FENOFIBRATE

NORTH

AB		145MG	A090715 002	Apr 05, 2012
AB	YICHANG HUMANWELL	54MG	A218548 001	Apr 24, 2024
AB		160MG	A218548 002	Apr 24, 2024
BX	CREEKWOOD PHARMS	40MG	A217732 001	Sep 07, 2023
BX		120MG	A217732 002	Sep 07, 2023
	SUN PHARM INDS LTD	107MG	A076635 002	Oct 31, 2005

FENOPROFEN CALCIUM

CAPSULE; ORAL

FENOPROFEN CALCIUM

RISING

		EQ 200MG BASE	A214475 002	Jul 26, 2024
		EQ 300MG BASE	A214475 003	Jul 26, 2024
!		EQ 400MG BASE	A214475 001	Jul 18, 2022

FENTANYL

FILM, EXTENDED RELEASE; TRANSDERMAL

FENTANYL-100

AB	DIFGEN PHARMS	100MCG/HR	A077449 004	Oct 20, 2008
AB	KINDEVA	100MCG/HR	A202097 005	Nov 04, 2016
AB	MYLAN TECHNOLOGIES	100MCG/HR	A076258 004	Jan 28, 2005
AB	SPECGX LLC	100MCG/HR	A077154 004	Feb 09, 2011

FENTANYL-12

AB	DIFGEN PHARMS	12.5MCG/HR	A077449 005	Sep 11, 2015
AB	KINDEVA	12.5MCG/HR	A202097 001	Nov 04, 2016
AB	MYLAN TECHNOLOGIES	12.5MCG/HR	A076258 005	Jan 23, 2007
AB	SPECGX LLC	12.5MCG/HR	A077154 005	Jun 11, 2015

FENTANYL-25

AB	DIFGEN PHARMS	25MCG/HR	A077449 001	Oct 20, 2008
AB	KINDEVA	25MCG/HR	A202097 002	Nov 04, 2016
AB	MYLAN TECHNOLOGIES	25MCG/HR	A076258 001	Jan 28, 2005
AB	! SPECGX LLC	25MCG/HR	A077154 001	Feb 09, 2011

FENTANYL-37

AB	DIFGEN PHARMS	37.5MCG/HR	A077449 006	Dec 06, 2017
AB	MYLAN TECHNOLOGIES	37.5MCG/HR	A076258 006	Dec 29, 2014
AB	SPECGX LLC	37.5MCG/HR	A077154 006	Jan 14, 2020

FENTANYL-50

AB	DIFGEN PHARMS	50MCG/HR	A077449 002	Oct 20, 2008
AB	KINDEVA	50MCG/HR	A202097 003	Nov 04, 2016
AB	MYLAN TECHNOLOGIES	50MCG/HR	A076258 002	Jan 28, 2005
AB	SPECGX LLC	50MCG/HR	A077154 002	Feb 09, 2011

FENTANYL-62

AB	DIFGEN PHARMS	62.5MCG/HR	A077449 007	Dec 06, 2017
AB	MYLAN TECHNOLOGIES	62.5MCG/HR	A076258 007	Dec 29, 2014
AB	SPECGX LLC	62.5MCG/HR	A077154 007	Jan 14, 2020

FENTANYL-75

AB	DIFGEN PHARMS	75MCG/HR	A077449 003	Oct 20, 2008
AB	KINDEVA	75MCG/HR	A202097 004	Nov 04, 2016
AB	MYLAN TECHNOLOGIES	75MCG/HR	A076258 003	Jan 28, 2005
AB	SPECGX LLC	75MCG/HR	A077154 003	Feb 09, 2011

FENTANYL-87

AB	DIFGEN PHARMS	87.5MCG/HR	A077449 008	Dec 06, 2017
AB	MYLAN TECHNOLOGIES	87.5MCG/HR	A076258 008	Dec 29, 2014

FENTANYL CITRATE

INJECTABLE; INJECTION

FENTANYL CITRATE

AP	FRESENIUS KABI USA	EQ 0.05MG BASE/ML	A212086 001	Sep 01, 2020
AP	+! HIKMA	EQ 0.05MG BASE/ML	N019101 001	Jul 11, 1984
AP	HOSPIRA	EQ 0.05MG BASE/ML	N019115 001	Jan 12, 1985

FENTANYL CITRATE PRESERVATIVE FREE

AP	FRESENIUS KABI USA	EQ 0.05MG BASE/ML	A210762 001	May 03, 2019
-----------	--------------------	--------------------------	--------------------	--------------

SUBLIMAZE PRESERVATIVE FREE

AP	+! RISING	EQ 0.05MG BASE/ML	N016619 001	
	FENTANYL CITRATE			
	+! HIKMA	EQ 0.025MG BASE/0.5ML	N019101 002	Jan 20, 2023

PRESCRIPTION DRUG PRODUCT LIST

FERRIC CARBOXYMALTOSE

SOLUTION; INTRAVENOUS

INJECTAFER

+	!	AM REGENT	750MG IRON/15ML (50MG IRON/ML)	N203565	001	Jul 25, 2013
+	!		100MG IRON/2ML (50MG IRON/ML)	N203565	004	Feb 04, 2022
+	!		500MG IRON/10ML (50MG IRON/ML)	N203565	002	Oct 08, 2020
+	!		1GM IRON/20ML (50MG IRON/ML)	N203565	003	Apr 28, 2021

FERRIC CITRATE

TABLET; ORAL

AURYXIA

+	!	KERYX BIOPHARMS	EQ 210MG IRON	N205874	001	Sep 05, 2014
---	---	-----------------	---------------	---------	-----	--------------

FERRIC DERISOMALTOSE

SOLUTION; INTRAVENOUS

MONOFERRIC

+	!	PHARMACOSMOS	1GM/10ML (100MG/ML)	N208171	003	Jan 16, 2020
---	---	--------------	---------------------	---------	-----	--------------

FERRIC HEXACYANOFERRATE (II)

CAPSULE; ORAL

RADIOGARDASE (PRUSSIAN BLUE)

+	!	HEYL CHEMISCH	500MG	N021626	001	Oct 02, 2003
---	---	---------------	-------	---------	-----	--------------

FERRIC MALTOL

CAPSULE; ORAL

ACCRUFER

+	!	SHIELD TX	30MG IRON	N212320	001	Jul 25, 2019
---	---	-----------	-----------	---------	-----	--------------

FERRIC OXYHYDROXIDE

INJECTABLE; INJECTION

FERRLECIT

AB	+	!	SANOFI AVENTIS US	<u>EQ 62.5MG IRON/5ML (EQ 12.5MG IRON/ML)</u>	<u>N020955</u>	<u>001</u>	Feb 18, 1999
-----------	---	---	-------------------	--	-----------------------	-------------------	--------------

SODIUM FERRIC GLUCONATE COMPLEX IN SUCROSE

AB			HIKMA	<u>EQ 62.5MG IRON/5ML (EQ 12.5MG IRON/ML)</u>	<u>A078215</u>	<u>001</u>	Mar 31, 2011
-----------	--	--	-------	--	-----------------------	-------------------	--------------

INFED

BP	+	!	ALLERGAN	EQ 100MG IRON/2ML (EQ 50MG IRON/ML)	N017441	001	
----	---	---	----------	-------------------------------------	---------	-----	--

INJECTABLE; INTRAVENOUS

VENOFER

+			AM REGENT	EQ 50MG IRON/2.5ML (EQ 20MG IRON/ML)	N021135	002	Mar 20, 2005
+	!			EQ 100MG IRON/5ML (EQ 20MG IRON/ML)	N021135	001	Nov 06, 2000
+				EQ 200MG IRON/10ML (EQ 20MG IRON/ML)	N021135	004	Feb 09, 2007

TABLET, CHEWABLE; ORAL

VELPHORO

+	!	VIFOR FRESENIUS	EQ 500MG IRON	N205109	001	Nov 27, 2013
---	---	-----------------	---------------	---------	-----	--------------

FERUMOXYTOL

SOLUTION; INTRAVENOUS

FERAHEME

AB	+	!	COVIS	<u>EQ 510MG IRON/17ML (EQ 30MG IRON/ML)</u>	<u>N022180</u>	<u>001</u>	Jun 30, 2009
-----------	---	---	-------	--	-----------------------	-------------------	--------------

FERUMOXYTOL

AB			SANDOZ	<u>EQ 510MG IRON/17ML (EQ 30MG IRON/ML)</u>	<u>A206604</u>	<u>001</u>	Jan 15, 2021
-----------	--	--	--------	--	-----------------------	-------------------	--------------

FESOTERODINE FUMARATE

TABLET, EXTENDED RELEASE; ORAL

FESOTERODINE FUMARATE

AB			ALEMBIC	<u>4MG</u>	<u>A204973</u>	<u>001</u>	Jan 04, 2023
-----------	--	--	---------	-------------------	-----------------------	-------------------	--------------

AB				<u>8MG</u>	<u>A204973</u>	<u>002</u>	Jan 04, 2023
-----------	--	--	--	-------------------	-----------------------	-------------------	--------------

AB			ALKEM LABS LTD	<u>4MG</u>	<u>A204827</u>	<u>001</u>	Dec 10, 2015
-----------	--	--	----------------	-------------------	-----------------------	-------------------	--------------

AB				<u>8MG</u>	<u>A204827</u>	<u>002</u>	Dec 10, 2015
-----------	--	--	--	-------------------	-----------------------	-------------------	--------------

AB			AMNEAL PHARMS NY	<u>4MG</u>	<u>A205002</u>	<u>001</u>	Jan 04, 2023
-----------	--	--	------------------	-------------------	-----------------------	-------------------	--------------

AB				<u>8MG</u>	<u>A205002</u>	<u>002</u>	Jan 04, 2023
-----------	--	--	--	-------------------	-----------------------	-------------------	--------------

AB			AUROBINDO PHARMA	<u>4MG</u>	<u>A205007</u>	<u>001</u>	Feb 17, 2017
-----------	--	--	------------------	-------------------	-----------------------	-------------------	--------------

AB				<u>8MG</u>	<u>A205007</u>	<u>002</u>	Feb 17, 2017
-----------	--	--	--	-------------------	-----------------------	-------------------	--------------

AB			DR REDDYS	<u>4MG</u>	<u>A204975</u>	<u>001</u>	Aug 13, 2019
-----------	--	--	-----------	-------------------	-----------------------	-------------------	--------------

AB				<u>8MG</u>	<u>A204975</u>	<u>002</u>	Aug 13, 2019
-----------	--	--	--	-------------------	-----------------------	-------------------	--------------

AB			HETERO LABS LTD V	<u>4MG</u>	<u>A204792</u>	<u>001</u>	Jan 09, 2024
-----------	--	--	-------------------	-------------------	-----------------------	-------------------	--------------

AB				<u>8MG</u>	<u>A204792</u>	<u>002</u>	Jan 09, 2024
-----------	--	--	--	-------------------	-----------------------	-------------------	--------------

AB			ZYDUS PHARMS	<u>4MG</u>	<u>A204946</u>	<u>001</u>	Oct 03, 2017
-----------	--	--	--------------	-------------------	-----------------------	-------------------	--------------

AB				<u>8MG</u>	<u>A204946</u>	<u>002</u>	Oct 03, 2017
-----------	--	--	--	-------------------	-----------------------	-------------------	--------------

TOVIAZ

AB	+		PFIZER	<u>4MG</u>	<u>N022030</u>	<u>001</u>	Oct 31, 2008
-----------	---	--	--------	-------------------	-----------------------	-------------------	--------------

AB	+	!		<u>8MG</u>	<u>N022030</u>	<u>002</u>	Oct 31, 2008
-----------	---	---	--	-------------------	-----------------------	-------------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

FEXINIDAZOLE

TABLET; ORAL

FEXINIDAZOLE

+! SANOFI

600MG

N214429 001 Jul 16, 2021

FEXOFENADINE HYDROCHLORIDE

TABLET; ORAL

FEXOFENADINE HYDROCHLORIDEAB DR REDDYS LABS LTD30MGA076502 001 Apr 11, 2006AB60MGA076502 002 Apr 11, 2006AB180MGA076502 003 Apr 11, 2006AB RISING60MGA077081 003 Apr 11, 2008AB180MGA077081 001 Apr 16, 2007AB TEVA30MGA076447 001 Sep 01, 2005AB60MGA076447 002 Sep 01, 2005AB180MGA076447 003 Sep 01, 2005FEZOLINETANT

TABLET; ORAL

VEOZAH

+! ASTELLAS

45MG

N216578 001 May 12, 2023

FIDAXOMICIN

FOR SUSPENSION; ORAL

DIFICID

+! CUBIST PHARMS LLC

40MG/ML

N213138 001 Jan 24, 2020

TABLET; ORAL

DIFICID

+! CUBIST PHARMS LLC

200MG

N201699 001 May 27, 2011

FINASTERIDE

TABLET; ORAL

FINASTERIDEAB ACCORD HLTHCARE1MGA091643 001 Nov 05, 2013AB5MGA090121 001 Feb 23, 2010AB ALKEM LABS LTD1MGA207750 001 Jan 06, 2017AB5MGA204304 001 Jan 05, 2017AB AUROBINDO PHARMA5MGA078341 001 Oct 30, 2007AB AUROBINDO PHARMA1MGA203687 001 Nov 05, 2013

LTD

AB DR REDDYS LABS INC1MGA076436 001 Jul 28, 2006AB DR REDDYS LABS LTD5MGA076437 001 Feb 28, 2007AB HETERO LABS LTD III1MGA090060 001 Jul 01, 2013AB5MGA090061 001 Jun 07, 2010AB SUN PHARM1MGA090508 001 Jul 01, 2013AB5MGA090507 001 Aug 16, 2011AB TEVA5MGA076511 001 Dec 15, 2006AB ZYDUS PHARMS USA5MGA078900 001 Dec 28, 2009

INC

PROPECIAAB +! ORGANON1MGN020788 001 Dec 19, 1997PROSCARAB +! ORGANON5MGN020180 001 Jun 19, 1992FINERENONE

TABLET; ORAL

KERENDIA

+ BAYER HLTHCARE

10MG

N215341 001 Jul 09, 2021

+!

20MG

N215341 002 Jul 09, 2021

FINGOLIMOD HYDROCHLORIDE

CAPSULE; ORAL

FINGOLIMOD HYDROCHLORIDEAB ACCORD HLTHCAREEQ 0.5MG BASEA207991 001 Oct 28, 2020AB ALKEM LABS LTDEQ 0.5MG BASEA208004 001 Dec 30, 2020AB APOTEXEQ 0.5MG BASEA207993 001 Dec 18, 2020AB AUROBINDO PHARMAEQ 0.5MG BASEA207983 001 Feb 28, 2024

LTD

AB BIOCON LTDEQ 0.5MG BASEA207979 001 Dec 04, 2019AB BIONPHARMAEQ 0.5MG BASEA210252 001 May 24, 2023AB DR REDDYSEQ 0.5MG BASEA208000 001 Mar 05, 2021AB EZRA VENTURESEQ 0.5MG BASEA207945 001 Dec 06, 2023AB GLENMARK PHARMS LTDEQ 0.5MG BASEA207985 001 Jun 18, 2020AB HEC PHARM CO LTDEQ 0.5MG BASEA207939 001 Nov 10, 2021AB HETERO LABS LTD VEQ 0.5MG BASEA207933 001 May 18, 2020AB MYLANEQ 0.5MG BASEA208005 001 Jan 19, 2021

PRESCRIPTION DRUG PRODUCT LISTFINGOLIMOD HYDROCHLORIDE

CAPSULE;ORAL

FINGOLIMOD HYDROCHLORIDE

AB	PRINSTON INC	EQ 0.5MG BASE	A208003 001	Sep 07, 2022
AB	TEVA PHARMS USA	EQ 0.5MG BASE	A208008 001	Jul 02, 2020
AB	ZYDUS PHARMS	EQ 0.5MG BASE	A207994 001	Oct 14, 2020
GILENYA				
AB	+! NOVARTIS	EQ 0.5MG BASE	N022527 001	Sep 21, 2010
	+	EQ 0.25MG BASE	N022527 002	May 11, 2018

FINGOLIMOD LAURYL SULFATE

TABLET, ORALLY DISINTEGRATING;ORAL

TASCENSO ODT

	+ CYCLE	EQ 0.25MG BASE	N214962 001	Dec 23, 2021
	+!	EQ 0.5MG BASE	N214962 002	Dec 09, 2022

FISH OIL TRIGLYCERIDES

EMULSION;INTRAVENOUS

OMEGAVEN

	+! FRESENIUS KABI USA	5GM/50ML (0.1GM/ML)	N210589 001	Jul 27, 2018
	+!	10GM/100ML (0.1GM/ML)	N210589 002	Jul 27, 2018

FISH OIL; MEDIUM CHAIN TRIGLYCERIDES; OLIVE OIL; SOYBEAN OIL

EMULSION;INTRAVENOUS

SMOFLIPID 20%

	+! FRESENIUS KABI USA	3GM/100ML;6GM/100ML;5GM/100ML;6GM/100ML (100ML)	N207648 001	Jul 13, 2016
	+!	3GM/100ML;6GM/100ML;5GM/100ML;6GM/100ML (250ML)	N207648 002	Jul 13, 2016
	+!	3GM/100ML;6GM/100ML;5GM/100ML;6GM/100ML (500ML)	N207648 003	Jul 13, 2016
	+!	3GM/100ML;6GM/100ML;5GM/100ML;6GM/100ML (1000ML)	N207648 004	Aug 10, 2018

FLAVOXATE HYDROCHLORIDE

TABLET;ORAL

FLAVOXATE HYDROCHLORIDE

AB	EPIC PHARMA	100MG	A076835 001	Nov 30, 2005
AB	! PADAGIS US	100MG	A076831 001	Dec 16, 2004

FLECAINIDE ACETATE

TABLET;ORAL

FLECAINIDE ACETATE

AB	AMNEAL PHARM	50MG	A075442 001	Jul 31, 2001
AB		100MG	A075442 002	Jul 31, 2001
AB		150MG	A075442 003	Jul 31, 2001
AB	ANI PHARMS	50MG	A075882 001	Oct 28, 2002
AB		100MG	A075882 002	Oct 28, 2002
AB		150MG	A075882 003	Oct 28, 2002
AB	AUROBINDO PHARMA LTD	50MG	A202821 001	Nov 03, 2017
AB		100MG	A202821 002	Nov 03, 2017
AB		150MG	A202821 003	Nov 03, 2017
AB	BEXIMCO PHARMS USA	50MG	A210683 001	Sep 16, 2020
AB		100MG	A210683 002	Sep 16, 2020
AB		150MG	A210683 003	Sep 16, 2020
AB	HIKMA	50MG	A076278 001	Jan 14, 2003
AB		100MG	A076278 002	Jan 14, 2003
AB	!	150MG	A076278 003	Jan 14, 2003
AB	SUN PHARM INDS LTD	50MG	A076421 001	Mar 28, 2003
AB		100MG	A076421 002	Mar 28, 2003
AB		150MG	A076421 003	Mar 28, 2003
AB	YICHANG HUMANWELL	50MG	A215599 001	Sep 08, 2022
AB		100MG	A215599 002	Sep 08, 2022
AB		150MG	A215599 003	Sep 08, 2022

FLIBANSERIN

TABLET;ORAL

ADDYI

	+! SPROUT PHARMS	100MG	N022526 001	Aug 18, 2015
--	------------------	-------	-------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

FLORBETABEN F-18

SOLUTION; INTRAVENOUS

NEURACEQ

+! LIFE MOLECULAR 1.4-135mCi/ML N204677 001 Mar 19, 2014

FLORBETAPIR F-18

SOLUTION; INTRAVENOUS

AMYVID

+! AVID RADIOPHARMS 10-100ML (13.5-51mCi/ML) N202008 004 Oct 13, 2023

+! INC 10-50ML (13.5-51mCi/ML) N202008 003 Apr 06, 2012

FLORTAUCIPIR F-18

SOLUTION; INTRAVENOUS

TAUVID

+! AVID RADIOPHARMS 50ML (8.1-100mCi/ML) N212123 003 Jul 01, 2022

+! INC 100ML (8.1-100mCi/ML) N212123 004 Jul 01, 2022

FLOTUFOLASTAT F-18 GALLIUM

SOLUTION; INTRAVENOUS

POSLUMA

+! BLUE EARTH 25ML (8-158mCi/ML) N216023 001 May 25, 2023

FLOXURIDINE

INJECTABLE; INJECTION

FLOXURIDINE**AP** + FRESENIUS KABI USA **500MG/VIAL** **A075837 001** Feb 22, 2001**AP** ! HIKMA **500MG/VIAL** **A075387 001** Apr 16, 2000FLUCICLOVINE F-18

SOLUTION; INTRAVENOUS

AXUMIN

+! BLUE EARTH 9-221mCi/ML N208054 001 May 27, 2016

FLUCONAZOLE

FOR SUSPENSION; ORAL

DIFLUCAN**AB** + PFIZER **50MG/5ML** **N020090 001** Dec 23, 1993**AB** +! **200MG/5ML** **N020090 002** Dec 23, 1993**FLUCONAZOLE****AB** AUROBINDO PHARMA **50MG/5ML** **A079150 001** Sep 18, 2009**AB** **200MG/5ML** **A079150 002** Sep 18, 2009**AB** ZHEJIANG POLY PHARM **200MG/5ML** **A215738 001** Sep 21, 2023

INJECTABLE; INJECTION

FLUCONAZOLE IN SODIUM CHLORIDE 0.9%**AP** FRESENIUS KABI USA **200MG/100ML (2MG/ML)** **A076145 001** Jul 29, 2004**AP** **400MG/200ML (2MG/ML)** **A076145 002** Jul 29, 2004**AP** HIKMA **200MG/100ML (2MG/ML)** **A076087 001** Jul 29, 2004**AP** **400MG/200ML (2MG/ML)** **A076087 003** Jul 29, 2004**AP** HIKMA FARMACEUTICA **200MG/100ML (2MG/ML)** **A076736 001** Aug 23, 2005**FLUCONAZOLE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER****AP** BAXTER HLTHCARE **200MG/100ML (2MG/ML)** **A076766 001** Jul 29, 2004**AP** **400MG/200ML (2MG/ML)** **A076766 002** Jul 29, 2004**AP** HIKMA **200MG/100ML (2MG/ML)** **A078107 001** Jul 30, 2008**AP** **400MG/200ML (2MG/ML)** **A078107 002** Jul 30, 2008**AP** HIKMA FARMACEUTICA **200MG/100ML (2MG/ML)** **A078698 001** Jan 30, 2012**AP** **400MG/200ML (2MG/ML)** **A078698 002** Jan 30, 2012**AP** HOSPIRA **200MG/100ML (2MG/ML)** **A076303 001** Jul 29, 2004**AP** **400MG/200ML (2MG/ML)** **A076303 002** Jul 29, 2004**AP** ! INFORLIFE **200MG/100ML (2MG/ML)** **A079104 001** Jul 30, 2009**AP** ! **400MG/200ML (2MG/ML)** **A079104 002** Jul 30, 2009**AP** WOODWARD **200MG/100ML (2MG/ML)** **A077909 001** May 26, 2010**AP** **400MG/200ML (2MG/ML)** **A077909 002** May 26, 2010

FLUCONAZOLE IN SODIUM CHLORIDE 0.9%

HIKMA 100MG/50ML (2MG/ML) A076087 002 Sep 26, 2008

FLUCONAZOLE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

WOODWARD 100MG/50ML (2MG/ML) A077909 003 Apr 20, 2015

TABLET; ORAL

DIFLUCAN**AB** + PFIZER **50MG** **N019949 001** Jan 29, 1990**AB** + **100MG** **N019949 002** Jan 29, 1990**AB** + **150MG** **N019949 004** Jun 30, 1994**AB** +! **200MG** **N019949 003** Jan 29, 1990

PRESCRIPTION DRUG PRODUCT LIST

FLUCONAZOLE

TABLET; ORAL

FLUCONAZOLE

<u>AB</u>	ANI PHARMS	<u>50MG</u>	<u>A078423 001</u>	Mar 07, 2011
<u>AB</u>		<u>100MG</u>	<u>A078423 002</u>	Mar 07, 2011
<u>AB</u>		<u>150MG</u>	<u>A078423 003</u>	Mar 07, 2011
<u>AB</u>		<u>200MG</u>	<u>A078423 004</u>	Mar 07, 2011
<u>AB</u>	AUROBINDO PHARMA	<u>50MG</u>	<u>A077731 001</u>	Oct 07, 2008
<u>AB</u>		<u>100MG</u>	<u>A077731 002</u>	Oct 07, 2008
<u>AB</u>		<u>150MG</u>	<u>A077731 003</u>	Oct 07, 2008
<u>AB</u>		<u>200MG</u>	<u>A077731 004</u>	Oct 07, 2008
<u>AB</u>	CHARTWELL	<u>50MG</u>	<u>A076665 001</u>	Jul 29, 2004
<u>AB</u>		<u>100MG</u>	<u>A076665 002</u>	Jul 29, 2004
<u>AB</u>		<u>150MG</u>	<u>A076665 003</u>	Jul 29, 2004
<u>AB</u>		<u>200MG</u>	<u>A076665 004</u>	Jul 29, 2004
<u>AB</u>	DR REDDYS LABS INC	<u>50MG</u>	<u>A076658 001</u>	Jul 29, 2004
<u>AB</u>		<u>100MG</u>	<u>A076658 002</u>	Jul 29, 2004
<u>AB</u>		<u>150MG</u>	<u>A076658 003</u>	Jul 29, 2004
<u>AB</u>		<u>200MG</u>	<u>A076658 004</u>	Jul 29, 2004
<u>AB</u>	GLENMARK PHARMS LTD	<u>50MG</u>	<u>A077253 001</u>	Jan 25, 2006
<u>AB</u>		<u>100MG</u>	<u>A077253 002</u>	Jan 25, 2006
<u>AB</u>		<u>150MG</u>	<u>A077253 003</u>	Jan 25, 2006
<u>AB</u>		<u>200MG</u>	<u>A077253 004</u>	Jan 25, 2006
<u>AB</u>	TARO	<u>50MG</u>	<u>A076507 001</u>	Jul 29, 2004
<u>AB</u>		<u>100MG</u>	<u>A076507 002</u>	Jul 29, 2004
<u>AB</u>		<u>150MG</u>	<u>A076507 003</u>	Jul 29, 2004
<u>AB</u>		<u>200MG</u>	<u>A076507 004</u>	Jul 29, 2004
<u>AB</u>	THINQ PHARM-CRO PVT	<u>50MG</u>	<u>A076957 001</u>	Sep 28, 2005
<u>AB</u>		<u>100MG</u>	<u>A076957 002</u>	Sep 28, 2005
<u>AB</u>		<u>150MG</u>	<u>A076957 004</u>	Feb 27, 2017
<u>AB</u>		<u>200MG</u>	<u>A076957 003</u>	Sep 28, 2005
<u>AB</u>	ZYDUS PHARMS	<u>50MG</u>	<u>A208963 001</u>	Feb 16, 2017
<u>AB</u>		<u>100MG</u>	<u>A208963 002</u>	Feb 16, 2017
<u>AB</u>		<u>150MG</u>	<u>A208963 003</u>	Feb 16, 2017
<u>AB</u>		<u>200MG</u>	<u>A208963 004</u>	Feb 16, 2017

FLUCYDOSINE

CAPSULE; ORAL

ANCOBON

<u>AB</u>	+ BAUSCH	<u>250MG</u>	<u>N017001 001</u>	
<u>AB</u>	+!	<u>500MG</u>	<u>N017001 002</u>	

FLUCYDOSINE

<u>AB</u>	AUROBINDO PHARMA LTD	<u>250MG</u>	<u>A213665 001</u>	May 01, 2020
<u>AB</u>		<u>500MG</u>	<u>A213665 002</u>	May 01, 2020
<u>AB</u>	NOVEL LABS INC	<u>250MG</u>	<u>A204652 001</u>	Jul 07, 2017
<u>AB</u>		<u>500MG</u>	<u>A204652 002</u>	Jul 07, 2017
<u>AB</u>	SIGMAPHARM LABS LLC	<u>250MG</u>	<u>A201566 001</u>	Jun 28, 2011
<u>AB</u>		<u>500MG</u>	<u>A201566 002</u>	Jun 28, 2011
<u>AB</u>	STRIDES PHARMA	<u>250MG</u>	<u>A212632 001</u>	Apr 17, 2020
<u>AB</u>		<u>500MG</u>	<u>A212632 002</u>	Apr 17, 2020

FLUDARABINE PHOSPHATE

INJECTABLE; INJECTION

FLUDARABINE PHOSPHATE

<u>AP</u>	! ACTAVIS TOTOWA	<u>50MG/VIAL</u>	<u>A078610 001</u>	Feb 11, 2009
<u>AP</u>	FRESENIUS KABI USA	<u>50MG/VIAL</u>	<u>A078544 001</u>	Oct 15, 2007
<u>AP</u>	HIKMA	<u>50MG/VIAL</u>	<u>A076349 001</u>	Aug 28, 2003
<u>AP1</u>	AREVA PHARMS	<u>50MG/2ML (25MG/ML)</u>	<u>A090724 001</u>	Sep 27, 2010
<u>AP1</u>	FRESENIUS KABI USA	<u>50MG/2ML (25MG/ML)</u>	<u>A078393 001</u>	Oct 15, 2007
<u>AP1</u>	SAGENT PHARMS INC	<u>50MG/2ML (25MG/ML)</u>	<u>A076661 001</u>	Apr 28, 2004

FLUDEOXYGLUCOSE F-18

INJECTABLE; INTRAVENOUS

FLUDEOXYGLUCOSE F18

<u>AP</u>	3D IMAGING DRUG	<u>20-300mCi/ML</u>	<u>A203778 001</u>	Oct 30, 2015
<u>AP</u>	BIOMEDCL RES FDN	<u>20-300mCi/ML</u>	<u>A203710 001</u>	May 01, 2015
<u>AP</u>		<u>20-300mCi/ML</u>	<u>A203837 001</u>	May 01, 2015
<u>AP</u>	BRIGHAM WOMENS	<u>20-300mCi/ML</u>	<u>A203816 001</u>	Oct 30, 2014
<u>AP</u>	CARDINAL HEALTH 414	<u>20-300mCi/ML</u>	<u>A203603 001</u>	Nov 13, 2015
<u>AP</u>	!	<u>20-500mCi/ML</u>	<u>A203603 002</u>	Sep 27, 2018
<u>AP</u>	CHILDRENS HOSP MI	<u>20-300mCi/ML</u>	<u>A204385 001</u>	Oct 29, 2014
<u>AP</u>	DECATUR	<u>20-300mCi/ML</u>	<u>A204463 001</u>	Oct 21, 2014
<u>AP</u>	+! FEINSTEIN	<u>20-400mCi/ML</u>	<u>N021870 002</u>	Nov 21, 2008

PRESCRIPTION DRUG PRODUCT LIST

FLUDEOXYGLUCOSE F-18

INJECTABLE; INTRAVENOUS

FLUDEOXYGLUCOSE F18

<u>AP</u>	ISOLOGIC INNOVATIVE	<u>20-300mCi/ML</u>	<u>A204525</u>	<u>001</u>	Oct 29, 2014
<u>AP</u>	JUBILANT DRAXIMAGE	<u>20-300mCi/ML</u>	<u>A203920</u>	<u>001</u>	Jun 23, 2015
<u>AP</u>	KETTERING MEDCTR	<u>4-40mCi/ML</u>	<u>A204759</u>	<u>001</u>	Oct 27, 2015
<u>AP</u>	KREITCHMAN PET CTR	<u>10-100mCi/ML</u>	<u>A203942</u>	<u>001</u>	Apr 11, 2016
<u>AP</u>	MA GENERAL HOSP	<u>20-300mCi/ML</u>	<u>A204333</u>	<u>001</u>	Sep 25, 2014
<u>AP</u>	MCPRF	<u>20-240mCi/ML</u>	<u>A203612</u>	<u>001</u>	Aug 05, 2013
<u>AP</u>	MEM SLOAN-KETTERING	<u>20-300mCi/ML</u>	<u>A208679</u>	<u>001</u>	Dec 08, 2016
<u>AP</u>	METHODIST HOSP RES	<u>20-300mCi/ML</u>	<u>A203904</u>	<u>001</u>	Apr 23, 2015
<u>AP</u>	MIPS CRF	<u>20-300mCi/ML</u>	<u>A204472</u>	<u>001</u>	Sep 11, 2015
<u>AP</u>	NCM USA BRONX LLC	<u>20-300mCi/ML</u>	<u>A204512</u>	<u>001</u>	Jan 07, 2015
<u>AP</u>	! PETNET	<u>20-200mCi/ML</u>	<u>A079086</u>	<u>001</u>	Feb 25, 2011
<u>AP</u>	PHARMALOGIC	<u>20-300mCi/ML</u>	<u>A204264</u>	<u>001</u>	Dec 18, 2014
<u>AP</u>	PHARMALOGIC HLDGS	<u>20-200mCi/ML</u>	<u>A203664</u>	<u>001</u>	Feb 04, 2014
<u>AP</u>	PRECISION NUCLEAR	<u>20-500mCi/ML</u>	<u>A204546</u>	<u>001</u>	Apr 07, 2015
<u>AP</u>	! QUEEN HAMAMATSU PET	<u>10-100mCi/ML</u>	<u>A203771</u>	<u>001</u>	Aug 31, 2015
<u>AP</u>	SOFIE	<u>20-300mCi/ML</u>	<u>A203591</u>	<u>001</u>	Aug 31, 2015
<u>AP</u>	TRUSTEES UNIV PA	<u>20-200mCi/ML</u>	<u>A203801</u>	<u>001</u>	Oct 29, 2014
<u>AP</u>	! UCLA BIOMEDICAL	<u>4-40mCi/ML</u>	<u>A203811</u>	<u>001</u>	Jun 27, 2013
<u>AP</u>	UCSF RODIOPHARM	<u>20-300mCi/ML</u>	<u>A203902</u>	<u>001</u>	May 09, 2014
<u>AP</u>	UIHC PET IMAGING	<u>20-300mCi/ML</u>	<u>A203990</u>	<u>001</u>	Aug 06, 2014
<u>AP</u>	UNIV MICHIGAN	<u>20-300mCi/ML</u>	<u>A204531</u>	<u>001</u>	Jul 17, 2015
<u>AP</u>	UNIV SOUTHERN CA	<u>20-300mCi/ML</u>	<u>A209341</u>	<u>001</u>	Dec 16, 2020
<u>AP</u>	UNIV TX MD ANDERSON	<u>20-300mCi/ML</u>	<u>A203246</u>	<u>002</u>	Jan 13, 2014
<u>AP</u>	UNIV UTAH CYCLOTRON	<u>20-300mCi/ML</u>	<u>A204498</u>	<u>001</u>	Jun 23, 2015
<u>AP</u>	WISCONSIN	<u>20-500mCi/ML</u>	<u>A203709</u>	<u>001</u>	Oct 23, 2013
<u>AP</u>	WUSM CYCLOTRON	<u>20-300mCi/ML</u>	<u>A203935</u>	<u>001</u>	Feb 05, 2014
	NORTHLAND	4-500mCi/ML	A203994	001	Feb 04, 2015
	NUKEMED	4-500mCi/ML	A203911	001	Apr 22, 2015
	UNIV TX MD ANDERSON	20-150mCi/ML	A203246	001	Jan 13, 2014

FLUDROCORTISONE ACETATE

TABLET; ORAL

FLUDROCORTISONE ACETATE

<u>AB</u>	BARR	<u>0.1MG</u>	<u>A040425</u>	<u>001</u>	Jan 21, 2003
<u>AB</u>	! IMPAX LABS	<u>0.1MG</u>	<u>A040431</u>	<u>001</u>	Mar 18, 2002
<u>AB</u>	NOVITIUM PHARMA	<u>0.1MG</u>	<u>A215279</u>	<u>001</u>	May 31, 2022
<u>AB</u>	ZYDUS LIFESCIENCES	<u>0.1MG</u>	<u>A219251</u>	<u>001</u>	Oct 16, 2024

FLUMAZENIL

INJECTABLE; INJECTION

FLUMAZENIL

<u>AP</u>	BAXTER HLTHCARE CORP	<u>0.5MG/5ML (0.1MG/ML)</u>	<u>A076755</u>	<u>002</u>	Oct 12, 2004
<u>AP</u>		<u>1MG/10ML (0.1MG/ML)</u>	<u>A076755</u>	<u>001</u>	Oct 12, 2004
<u>AP</u>	FRESENIUS KABI USA	<u>0.5MG/5ML (0.1MG/ML)</u>	<u>A076955</u>	<u>002</u>	Oct 12, 2004
<u>AP</u>		<u>1MG/10ML (0.1MG/ML)</u>	<u>A076955</u>	<u>001</u>	Oct 12, 2004
<u>AP</u>	HIKMA	<u>0.5MG/5ML (0.1MG/ML)</u>	<u>A076256</u>	<u>002</u>	Oct 12, 2004
<u>AP</u>		<u>0.5MG/5ML (0.1MG/ML)</u>	<u>A076787</u>	<u>002</u>	Oct 12, 2004
<u>AP</u>		<u>1MG/10ML (0.1MG/ML)</u>	<u>A076256</u>	<u>001</u>	Oct 12, 2004
<u>AP</u>		<u>1MG/10ML (0.1MG/ML)</u>	<u>A076787</u>	<u>001</u>	Oct 12, 2004
<u>AP</u>	HIKMA FARMACEUTICA	<u>0.5MG/5ML (0.1MG/ML)</u>	<u>A078527</u>	<u>001</u>	Mar 23, 2009
<u>AP</u>	!	<u>1MG/10ML (0.1MG/ML)</u>	<u>A078527</u>	<u>002</u>	Mar 23, 2009
<u>AP</u>	RISING	<u>0.5MG/5ML (0.1MG/ML)</u>	<u>A078595</u>	<u>001</u>	May 13, 2008
<u>AP</u>	SAGENT PHARMS	<u>0.5MG/5ML (0.1MG/ML)</u>	<u>A090584</u>	<u>001</u>	Aug 28, 2012
<u>AP</u>		<u>1MG/10ML (0.1MG/ML)</u>	<u>A090584</u>	<u>002</u>	Aug 28, 2012
<u>AP</u>	SANDOZ	<u>0.5MG/5ML (0.1MG/ML)</u>	<u>A077071</u>	<u>001</u>	May 03, 2005
<u>AP</u>		<u>1MG/10ML (0.1MG/ML)</u>	<u>A077071</u>	<u>002</u>	May 03, 2005

FLUNISOLIDE

SPRAY, METERED; NASAL

FLUNISOLIDE

<u>AB</u>	! BAUSCH	<u>0.025MG/SPRAY</u>	<u>A074805</u>	<u>001</u>	Feb 20, 2002
<u>AB</u>	RICONPHARMA LLC	<u>0.025MG/SPRAY</u>	<u>A207802</u>	<u>001</u>	Jun 16, 2022
<u>AB</u>	RISING	<u>0.025MG/SPRAY</u>	<u>A077704</u>	<u>001</u>	Aug 03, 2006

FLUOCINOLONE ACETONIDE

CREAM; TOPICAL

FLUOCINOLONE ACETONIDE

<u>AT</u>	COSETTE	<u>0.01%</u>	<u>A089526</u>	<u>001</u>	Jul 26, 1988
<u>AT</u>		<u>0.025%</u>	<u>A210747</u>	<u>001</u>	Nov 05, 2018
<u>AT</u>	FOUGERA PHARMS INC	<u>0.01%</u>	<u>A088170</u>	<u>001</u>	Dec 16, 1982
<u>AT</u>		<u>0.025%</u>	<u>A088169</u>	<u>001</u>	Dec 16, 1982

PRESCRIPTION DRUG PRODUCT LIST

FLUOCINOLONE ACETONIDE

CREAM; TOPICAL

FLUOCINOLONE ACETONIDE

AT	TARO	0.025%	A087104 001	Apr 27, 1982
-----------	------	---------------	--------------------	--------------

SYNALAR

AT	+	MEDIMETRIKS PHARMS	0.01%	N012787 004
-----------	---	--------------------	--------------	--------------------

AT	+		0.025%	N012787 002
-----------	---	--	---------------	--------------------

AT	+		0.025%	N012787 005
-----------	---	--	---------------	--------------------

IMPLANT; INTRAVITREAL

ILUVIEN

+	ALIMERA SCIENCES INC	0.19MG	N201923 001	Sep 26, 2014
---	----------------------	--------	-------------	--------------

RETISERT

+	BAUSCH AND LOMB	0.59MG	N021737 001	Apr 08, 2005
---	-----------------	--------	-------------	--------------

YUTIQ

+	ALIMERA SCIENCES INC	0.18MG	N210331 001	Oct 12, 2018
---	----------------------	--------	-------------	--------------

OIL; TOPICAL

DERMA-SMOOTH/FS

AT	+	HILL DERMAC	0.01%	N019452 001	Feb 03, 1988
-----------	---	-------------	--------------	--------------------	--------------

AT	+		0.01%	N019452 002	Nov 09, 2005
-----------	---	--	--------------	--------------------	--------------

FLUCINOLONE ACETONIDE

AT	GLENMARK PHARMS LTD	0.01%	A210556 001	Oct 25, 2018
-----------	---------------------	--------------	--------------------	--------------

FLUOCINOLONE ACETONIDE

AT	NOVEL LABS INC	0.01%	A207345 001	Jul 31, 2023
-----------	----------------	--------------	--------------------	--------------

AT		0.01%	A207347 001	Aug 07, 2023
-----------	--	--------------	--------------------	--------------

AT	PADAGIS ISRAEL	0.01%	A202847 001	Aug 09, 2013
-----------	----------------	--------------	--------------------	--------------

AT		0.01%	A202848 001	Aug 09, 2013
-----------	--	--------------	--------------------	--------------

AT	QUAGEN	0.01%	A212760 001	Apr 02, 2021
-----------	--------	--------------	--------------------	--------------

AT		0.01%	A212761 001	Apr 02, 2021
-----------	--	--------------	--------------------	--------------

AT	RISING	0.01%	A090982 001	Apr 25, 2016
-----------	--------	--------------	--------------------	--------------

AT		0.01%	A203377 001	Apr 25, 2016
-----------	--	--------------	--------------------	--------------

AT	TARO	0.01%	A202368 001	May 19, 2016
-----------	------	--------------	--------------------	--------------

AT		0.01%	A209336 001	May 19, 2016
-----------	--	--------------	--------------------	--------------

FLUOCINONIDE ACETONIDE

AT	GLENMARK PHARMS LTD	0.01%	A210539 001	Oct 26, 2018
-----------	---------------------	--------------	--------------------	--------------

OIL/DROPS; OTIC

DERMOTIC

AT	+	HILL DERMAC	0.01%	N019452 003	Nov 09, 2005
-----------	---	-------------	--------------	--------------------	--------------

FLAC

AT	ANDA REPOSITORY	0.01%	A210736 001	Apr 11, 2018
-----------	-----------------	--------------	--------------------	--------------

FLUOCINOLONE ACETONIDE

AT	PADAGIS ISRAEL	0.01%	A202849 001	Jul 17, 2017
-----------	----------------	--------------	--------------------	--------------

AT	QUAGEN	0.01%	A212762 001	Apr 02, 2021
-----------	--------	--------------	--------------------	--------------

AT	RISING	0.01%	A203378 001	Apr 25, 2016
-----------	--------	--------------	--------------------	--------------

AT	TRUPHARMA	0.01%	A213264 001	Feb 05, 2021
-----------	-----------	--------------	--------------------	--------------

FLUOCINONIDE ACETONIDE

AT	GLENMARK PHARMS LTD	0.01%	A211815 001	Dec 14, 2018
-----------	---------------------	--------------	--------------------	--------------

OINTMENT; TOPICAL

FLUOCINOLONE ACETONIDE

AT	COSETTE	0.025%	A089524 001	Jul 26, 1988
-----------	---------	---------------	--------------------	--------------

AT	FOUGERA PHARMS INC	0.025%	A088168 001	Dec 16, 1982
-----------	--------------------	---------------	--------------------	--------------

AT	TARO	0.025%	A040041 001	Sep 15, 1994
-----------	------	---------------	--------------------	--------------

SYNALAR

AT	+	MEDIMETRIKS PHARMS	0.025%	N013960 001
-----------	---	--------------------	---------------	--------------------

SOLUTION; TOPICAL

FLUOCINOLONE ACETONIDE

AT	CHARTWELL RX	0.01%	A209596 001	Dec 26, 2017
-----------	--------------	--------------	--------------------	--------------

AT	ENCUBE ETHICALS	0.01%	A209913 001	Feb 13, 2019
-----------	-----------------	--------------	--------------------	--------------

AT	FOUGERA PHARMS INC	0.01%	A088167 001	Dec 16, 1982
-----------	--------------------	--------------	--------------------	--------------

AT	LUPIN	0.01%	A206422 001	Sep 02, 2015
-----------	-------	--------------	--------------------	--------------

AT	TARO	0.01%	A089124 001	Sep 11, 1985
-----------	------	--------------	--------------------	--------------

SYNALAR

AT	+	MEDIMETRIKS PHARMS	0.01%	N015296 001
-----------	---	--------------------	--------------	--------------------

FLUOCINOLONE ACETONIDE; HYDROQUINONE; TRETINOIN

CREAM; TOPICAL

TRI-LUMA

+	GALDERMA LABS LP	0.01%; 4%; 0.05%	N021112 001	Jan 18, 2002
---	------------------	------------------	-------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

FLUOCINOLONE ACETONIDE; NEOMYCIN SULFATE

CREAM; TOPICAL

NEO-SYNALAR

! MEDIMETRIKS PHARMS 0.025%;EQ 3.5MG BASE/GM

A060700 001

FLUOCINONIDE

CREAM; TOPICAL

FLUOCINONIDE

AB	FOUGERA PHARMS INC	<u>0.1%</u>	<u>A200735</u>	<u>001</u>	Jul 14, 2014
AB	GLENMARK PHARMS LTD	<u>0.1%</u>	<u>A091282</u>	<u>001</u>	Jul 14, 2014
AB	TARO	<u>0.1%</u>	<u>A200734</u>	<u>001</u>	Jul 14, 2014
AB	ZYDUS LIFESCIENCES	<u>0.1%</u>	<u>A208989</u>	<u>001</u>	Feb 10, 2020

VANOS

AB	+! BAUSCH	<u>0.1%</u>	<u>N021758</u>	<u>001</u>	Feb 11, 2005
-----------	-----------	-------------	-----------------------	-------------------	--------------

FLUOCINONIDE

AB1	! AMNEAL	<u>0.05%</u>	<u>A210554</u>	<u>001</u>	Aug 21, 2018
AB1	COSETTE	<u>0.05%</u>	<u>A073085</u>	<u>001</u>	Feb 14, 1992
AB1	FOUGERA PHARMS INC	<u>0.05%</u>	<u>A073030</u>	<u>001</u>	Oct 17, 1994
AB1	TARO	<u>0.05%</u>	<u>A071500</u>	<u>001</u>	Jun 10, 1987
AB1	TEVA	<u>0.05%</u>	<u>A072488</u>	<u>001</u>	Feb 06, 1989

FLUOCINONIDE EMULSIFIED BASE

AB2	COSETTE	<u>0.05%</u>	<u>A074204</u>	<u>001</u>	Jun 13, 1995
AB2	FOUGERA PHARMS	<u>0.05%</u>	<u>A076586</u>	<u>001</u>	Jun 23, 2004
AB2	! TARO	<u>0.05%</u>	<u>A072494</u>	<u>001</u>	Jan 19, 1989
AB2	TEVA	<u>0.05%</u>	<u>A072490</u>	<u>001</u>	Feb 07, 1989

GEL; TOPICAL

FLUOCINONIDE

AB	FOUGERA PHARMS INC	<u>0.05%</u>	<u>A072933</u>	<u>001</u>	Dec 30, 1994
AB	! TARO	<u>0.05%</u>	<u>A074935</u>	<u>001</u>	Jul 29, 1997

OINTMENT; TOPICAL

FLUOCINONIDE

AB	CHARTWELL RX	<u>0.05%</u>	<u>A207538</u>	<u>001</u>	Jul 31, 2017
AB	FOUGERA PHARMS	<u>0.05%</u>	<u>A074905</u>	<u>001</u>	Aug 26, 1997
AB	! TARO	<u>0.05%</u>	<u>A075008</u>	<u>001</u>	Jun 30, 1999
AB	TEVA	<u>0.05%</u>	<u>A073481</u>	<u>001</u>	Dec 27, 1991
AB	XIROMED	<u>0.05%</u>	<u>A212976</u>	<u>001</u>	Nov 26, 2019

SOLUTION; TOPICAL

FLUOCINONIDE

AT	CHARTWELL RX	<u>0.05%</u>	<u>A209118</u>	<u>001</u>	Apr 23, 2018
AT	ENCUBE ETHICALS	<u>0.05%</u>	<u>A209699</u>	<u>001</u>	Nov 29, 2018
AT	FOUGERA PHARMS INC	<u>0.05%</u>	<u>A072934</u>	<u>001</u>	Feb 27, 1995
AT	MACLEODS PHARMS LTD	<u>0.05%</u>	<u>A209283</u>	<u>001</u>	Apr 23, 2018
AT	NOVEL LABS INC	<u>0.05%</u>	<u>A206003</u>	<u>001</u>	Jul 21, 2017
AT	! TARO	<u>0.05%</u>	<u>A074799</u>	<u>001</u>	Dec 31, 1996
AT	ZYDUS PHARMS	<u>0.05%</u>	<u>A208948</u>	<u>001</u>	Jul 17, 2018

FLUORESCEIN SODIUM

INJECTABLE; INTRAVENOUS

AK-FLUOR 10%

AP	+! LONG GROVE PHARMS	<u>EQ 500MG BASE/5ML (EQ 100MG BASE/ML)</u>	<u>N022186</u>	<u>001</u>	Aug 08, 2008
-----------	----------------------	---	-----------------------	-------------------	--------------

AK-FLUOR 25%

AP	+! LONG GROVE PHARMS	<u>EQ 500MG BASE/2ML (EQ 250MG BASE/ML)</u>	<u>N022186</u>	<u>002</u>	Aug 08, 2008
-----------	----------------------	---	-----------------------	-------------------	--------------

FLUORESCEIN SODIUM

AP	NEXUS PHARMS	<u>EQ 500MG BASE/5ML (EQ 100MG BASE/ML)</u>	<u>A215709</u>	<u>001</u>	Sep 25, 2023
AP		<u>EQ 500MG BASE/2ML (EQ 250MG BASE/ML)</u>	<u>A215709</u>	<u>002</u>	Sep 25, 2023

FLUORESCITE

AP	+! ALCON LABS INC	<u>EQ 500MG BASE/5ML (EQ 100MG BASE/ML)</u>	<u>N021980</u>	<u>001</u>	Mar 28, 2006
-----------	-------------------	---	-----------------------	-------------------	--------------

FLUORODOPA F-18

SOLUTION; INTRAVENOUS

FLUORODOPA F18

+! FEINSTEIN 0.42-8.33mCi/ML

N200655 001 Oct 10, 2019

FLUOROESTRADIOL F-18

SOLUTION; INTRAVENOUS

CERIANNA

+! GE HEALTHCARE 50ML (4-100mCi/ML)

N212155 001 May 20, 2020

FLUOROMETHOLONE

SUSPENSION/DROPS; OPHTHALMIC

FLUOROMETHOLONE

AB	AMNEAL	<u>0.1%</u>	<u>A216348</u>	<u>001</u>	Jan 09, 2024
-----------	--------	-------------	-----------------------	-------------------	--------------

FML

AB	+! ABBVIE	<u>0.1%</u>	<u>N016851</u>	<u>002</u>	Jul 28, 1982
-----------	-----------	-------------	-----------------------	-------------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

FLUOROMETHOLONE

SUSPENSION/DROPS;OPHTHALMIC

FML FORTE

+! ABBVIE

0.25%

N019216 001 Apr 23, 1986

FLUOROMETHOLONE ACETATE

SUSPENSION/DROPS;OPHTHALMIC

FLAREX

+! HARROW EYE

0.1%

N019079 001 Feb 11, 1986

FLUOROURACIL

CREAM;TOPICAL

EFUDEX**AB** +! EXTROVIS**5%****N016831 003**FLUOROURACIL**AB** ACCORD HLTHCARE**5%****A214845 001** Oct 07, 2021**AB** DR REDDYS LABS SA**5%****A077524 001** Apr 11, 2008**AB** TARO**5%****A090368 001** Mar 05, 2010

CARAC

+! EXTROVIS

0.5%

N020985 001 Oct 27, 2000

TOLAK

+! HILL DERMACEUTICALS

4%

N022259 001 Sep 18, 2015

INJECTABLE;INJECTION

FLUOROURACIL**AP** ! ACCORD HLTHCARE**500MG/10ML (50MG/ML)****A040743 002** Apr 26, 2007**AP** !**1GM/20ML (50MG/ML)****A040743 001** Apr 26, 2007**AP** !**2.5GM/50ML (50MG/ML)****A040798 002** Apr 26, 2007**AP** !**5GM/100ML (50MG/ML)****A040798 001** Apr 26, 2007**AP** ALEMBIC**2.5GM/50ML (50MG/ML)****A217295 001** Mar 03, 2023**AP****5GM/100ML (50MG/ML)****A217676 001** Oct 18, 2023**AP** EUGIA PHARMA**500MG/10ML (50MG/ML)****A202668 001** Jul 17, 2012

SPECLTS

AP**1GM/20ML (50MG/ML)****A202668 002** Jul 17, 2012**AP****2.5GM/50ML (50MG/ML)****A202669 001** Jul 17, 2012**AP****5GM/100ML (50MG/ML)****A202669 002** Jul 17, 2012**AP** FRESENIUS KABI USA**500MG/10ML (50MG/ML)****A040279 002** Sep 30, 1998**AP****1GM/20ML (50MG/ML)****A040279 001** Sep 30, 1998**AP****2.5GM/50ML (50MG/ML)****A040278 001** Sep 30, 1998**AP****5GM/100ML (50MG/ML)****A040278 002** Sep 30, 1998**AP** GLAND PHARMA LTD**500MG/10ML (50MG/ML)****A210123 001** Oct 27, 2017**AP****1GM/20ML (50MG/ML)****A210123 002** Oct 27, 2017**AP****2.5GM/50ML (50MG/ML)****A210124 001** Dec 26, 2017**AP****5GM/100ML (50MG/ML)****A210124 002** Dec 26, 2017**AP** KINDOS**2.5GM/50ML (50MG/ML)****A216494 001** Sep 24, 2024**AP** SAGENT PHARMS INC**500MG/10ML (50MG/ML)****A203608 001** May 11, 2017**AP****1GM/20ML (50MG/ML)****A203608 002** May 11, 2017**AP****2.5GM/50ML (50MG/ML)****A203609 001** Feb 17, 2016**AP****5GM/100ML (50MG/ML)****A203609 002** Feb 17, 2016

SOLUTION;TOPICAL

EFUDEX**AT** +! EXTROVIS**2%****N016831 001**FLUOROURACIL**AT** ENCUBE**5%****A215612 001** Nov 02, 2023**AT** TARO**2%****A076526 001** Nov 05, 2003**AT** !**5%****A076526 002** Nov 05, 2003FLUOXETINE HYDROCHLORIDE

CAPSULE;ORAL

FLUOXETINE HYDROCHLORIDE**AB** ALEMBIC PHARMS LTD**EQ 40MG BASE****A090223 003** Mar 19, 2009**AB** APNAR PHARMA LP**EQ 40MG BASE****A075049 003** Jan 29, 2002**AB** AUROBINDO PHARMA**EQ 40MG BASE****A078619 003** Jan 31, 2008**AB** CADILA PHARMS LTD**EQ 40MG BASE****A206993 003** May 23, 2019**AB** HERITAGE**EQ 40MG BASE****A201336 003** Oct 01, 2012**AB** IVAX SUB TEVA**EQ 40MG BASE****A075245 003** Sep 28, 2004

PHARMS

AB MARKSANS PHARMA**EQ 40MG BASE****A075465 003** Aug 02, 2001**AB** MICRO LABS**EQ 40MG BASE****A216232 003** Mar 29, 2022**AB** SCIEGEN PHARMS INC**EQ 40MG BASE****A204597 003** Mar 16, 2015**AB** STRIDES PHARMA**EQ 40MG BASE****A076922 003** Dec 16, 2004**AB** TEVA**EQ 40MG BASE****A075452 003** Jan 29, 2002PROZAC**AB** +! ELI LILLY AND CO**EQ 40MG BASE****N018936 003** Jun 15, 1999

PRESCRIPTION DRUG PRODUCT LIST

FLUOXETINE HYDROCHLORIDE

CAPSULE;ORAL

FLUOXETINE HYDROCHLORIDE

AB1	ALEMbic PHARMS LTD	EQ 10MG BASE	A090223 001	Mar 19, 2009
AB1		EQ 20MG BASE	A090223 002	Mar 19, 2009
AB1	APNAR PHARMA LP	EQ 10MG BASE	A075049 001	Aug 02, 2001
AB1		EQ 20MG BASE	A075049 002	Jan 29, 2002
AB1	AUROBINDO PHARMA	EQ 10MG BASE	A078619 001	Jan 31, 2008
AB1		EQ 20MG BASE	A078619 002	Jan 31, 2008
AB1	CADILA PHARMS LTD	EQ 10MG BASE	A206993 001	May 23, 2019
AB1		EQ 20MG BASE	A206993 002	May 23, 2019
AB1	HERITAGE	EQ 10MG BASE	A201336 001	Oct 01, 2012
AB1		EQ 20MG BASE	A201336 002	Oct 01, 2012
AB1	IVAX SUB TEVA PHARMS	EQ 10MG BASE	A075245 002	Jan 31, 2002
AB1		EQ 20MG BASE	A075245 001	Jan 31, 2002
AB1	LANDELA PHARM	EQ 10MG BASE	A075464 001	Jan 30, 2002
AB1		EQ 20MG BASE	A075464 002	Jan 30, 2002
AB1	MARKSANS PHARMA	EQ 10MG BASE	A075465 001	Jan 29, 2002
AB1		EQ 20MG BASE	A075465 002	Jan 29, 2002
AB1	MICRO LABS	EQ 10MG BASE	A216232 001	Mar 29, 2022
AB1		EQ 20MG BASE	A216232 002	Mar 29, 2022
AB1	SCIEGEN PHARMS INC	EQ 10MG BASE	A204597 001	Mar 16, 2015
AB1		EQ 20MG BASE	A204597 002	Mar 16, 2015
AB1	STRIDES PHARMA	EQ 10MG BASE	A076922 001	Dec 16, 2004
AB1		EQ 20MG BASE	A076922 002	Dec 16, 2004
AB1	TEVA	EQ 10MG BASE	A075452 001	Jan 29, 2002
AB1		EQ 20MG BASE	A075452 002	Jan 29, 2002
AB1	TEVA PHARMS USA	EQ 10MG BASE	A076001 001	Jan 29, 2002
AB1		EQ 20MG BASE	A076001 002	Jan 29, 2002

PROZAC

AB1	+	ELI LILLY AND CO	EQ 10MG BASE	N018936 006	Dec 23, 1992
AB1	+		EQ 20MG BASE	N018936 001	Dec 29, 1987

CAPSULE, DELAYED REL PELLETS;ORAL

FLUOXETINE HYDROCHLORIDE

! DR REDDYS LABS LTD EQ 90MG BASE

A078572 001 Mar 22, 2010

SOLUTION;ORAL

FLUOXETINE HYDROCHLORIDE

AA	AUROBINDO PHARMA	EQ 20MG BASE/5ML	A079209 001	Mar 20, 2009
AA	LANNETT CO INC	EQ 20MG BASE/5ML	A077849 001	Feb 09, 2007
AA	NOSTRUM LABS INC	EQ 20MG BASE/5ML	A075292 001	Feb 07, 2002
AA	NOVITIUM PHARMA	EQ 20MG BASE/5ML	A216448 001	Nov 09, 2022
AA	! PHARM ASSOC	EQ 20MG BASE/5ML	A076015 001	Jan 30, 2002
AA	TEVA	EQ 20MG BASE/5ML	A075506 001	Aug 02, 2001
AA	UPSHER SMITH LABS	EQ 20MG BASE/5ML	A216953 001	Nov 15, 2022

TABLET;ORAL

FLUOXETINE HYDROCHLORIDE

AB	ALEMbic	EQ 10MG BASE	A208698 001	Apr 05, 2017	
AB		EQ 20MG BASE	A208698 002	Apr 05, 2017	
AB	AUROBINDO PHARMA LTD	EQ 10MG BASE	A213286 001	Apr 08, 2020	
AB		EQ 20MG BASE	A213286 002	Apr 08, 2020	
AB		EQ 60MG BASE	A213265 001	Jun 10, 2020	
AB	DR REDDYS	EQ 10MG BASE	A076006 001	Jan 30, 2002	
AB	!	EQ 20MG BASE	A076006 002	Apr 23, 2018	
AB		EQ 60MG BASE	A211721 001	Jan 25, 2019	
AB	ENDO OPERATIONS	EQ 10MG BASE	A203836 001	Aug 19, 2016	
AB		EQ 20MG BASE	A203836 002	Aug 19, 2016	
AB		EQ 60MG BASE	A209419 001	Nov 16, 2017	
AB	INVENTIA HLTHCARE	EQ 60MG BASE	A209695 001	Nov 20, 2017	
AB	LUPIN LTD	EQ 10MG BASE	A211653 001	Apr 15, 2019	
AB		EQ 20MG BASE	A211653 002	Apr 15, 2019	
AB		EQ 60MG BASE	A211632 001	Feb 08, 2019	
AB	SCIEGEN PHARMS INC	EQ 10MG BASE	A210935 001	Mar 20, 2019	
AB		EQ 20MG BASE	A210935 002	Mar 20, 2019	
AB		EQ 60MG BASE	A211282 001	Jan 10, 2019	
AB	STRIDES PHARMA	EQ 10MG BASE	A212684 001	Apr 05, 2024	
AB		EQ 20MG BASE	A212684 002	Apr 05, 2024	
AB		EQ 60MG BASE	A212683 001	Sep 13, 2024	
AB	TARO	EQ 60MG BASE	A211477 001	Nov 21, 2018	
AB	+!	TWI PHARMS	EQ 60MG BASE	N202133 001	Oct 06, 2011
		TORRENT	EQ 10MG BASE	A206937 001	Oct 21, 2016
	!		EQ 20MG BASE	A206937 002	Oct 21, 2016

PRESCRIPTION DRUG PRODUCT LIST

FLUOXETINE HYDROCHLORIDE; OLANZAPINE

CAPSULE; ORAL

OLANZAPINE AND FLUOXETINE HYDROCHLORIDE

<u>AB</u>	ENDO OPERATIONS	<u>EQ 25MG BASE;EQ 3MG BASE</u>	<u>A077742 001</u>	Nov 02, 2012
<u>AB</u>		<u>EQ 25MG BASE;EQ 6MG BASE</u>	<u>A077742 002</u>	Nov 02, 2012
<u>AB</u>		<u>EQ 25MG BASE;EQ 12MG BASE</u>	<u>A077742 003</u>	Nov 02, 2012
<u>AB</u>		<u>EQ 50MG BASE;EQ 6MG BASE</u>	<u>A077742 004</u>	Nov 02, 2012
<u>AB</u>		<u>EQ 50MG BASE;EQ 12MG BASE</u>	<u>A077742 005</u>	Nov 02, 2012
<u>AB</u>	EPIC PHARMA LLC	<u>EQ 25MG BASE;EQ 3MG BASE</u>	<u>A078901 005</u>	Nov 16, 2012
<u>AB</u>		<u>EQ 25MG BASE;EQ 6MG BASE</u>	<u>A078901 001</u>	Nov 16, 2012
<u>AB</u>		<u>EQ 25MG BASE;EQ 12MG BASE</u>	<u>A078901 003</u>	Nov 16, 2012
<u>AB</u>		<u>EQ 50MG BASE;EQ 6MG BASE</u>	<u>A078901 002</u>	Nov 16, 2012
<u>AB</u>		<u>EQ 50MG BASE;EQ 12MG BASE</u>	<u>A078901 004</u>	Nov 16, 2012
<u>AB</u>	TEVA PHARMS	<u>EQ 25MG BASE;EQ 3MG BASE</u>	<u>A202074 001</u>	Mar 25, 2013
<u>AB</u>		<u>EQ 25MG BASE;EQ 6MG BASE</u>	<u>A077528 001</u>	Jun 19, 2012
<u>AB</u>		<u>EQ 25MG BASE;EQ 12MG BASE</u>	<u>A077528 002</u>	Jun 19, 2012
<u>AB</u>		<u>EQ 50MG BASE;EQ 6MG BASE</u>	<u>A077528 003</u>	Jun 19, 2012
<u>AB</u>		<u>EQ 50MG BASE;EQ 12MG BASE</u>	<u>A077528 004</u>	Jun 19, 2012
<u>SYMBYAX</u>				
<u>AB</u>	+ LILLY	<u>EQ 25MG BASE;EQ 3MG BASE</u>	<u>N021520 001</u>	Apr 09, 2007
<u>AB</u>	+	<u>EQ 25MG BASE;EQ 6MG BASE</u>	<u>N021520 002</u>	Dec 24, 2003
<u>AB</u>	+	<u>EQ 25MG BASE;EQ 12MG BASE</u>	<u>N021520 004</u>	Dec 24, 2003
<u>AB</u>	+!	<u>EQ 50MG BASE;EQ 6MG BASE</u>	<u>N021520 003</u>	Dec 24, 2003
<u>AB</u>	+	<u>EQ 50MG BASE;EQ 12MG BASE</u>	<u>N021520 005</u>	Dec 24, 2003

FLUPHENAZINE DECANOATE

INJECTABLE; INJECTION

FLUPHENAZINE DECANOATE

<u>AO</u>	ENDO OPERATIONS	<u>25MG/ML</u>	<u>A203732 001</u>	Jul 03, 2014
<u>AO</u>	EUGIA PHARMA	<u>25MG/ML</u>	<u>A207739 001</u>	Oct 17, 2017
<u>AO</u>	! FRESENIUS KABI USA	<u>25MG/ML</u>	<u>A071413 001</u>	Jul 14, 1987
<u>AO</u>	GLAND PHARMA LTD	<u>25MG/ML</u>	<u>A215509 001</u>	Mar 30, 2023
<u>AO</u>	HIKMA	<u>25MG/ML</u>	<u>A074531 001</u>	Aug 30, 1996
<u>AO</u>	MSN	<u>25MG/ML</u>	<u>A215365 001</u>	Nov 02, 2023
<u>AO</u>	MYLAN LABS LTD	<u>25MG/ML</u>	<u>A075918 001</u>	Aug 17, 2001

FLUPHENAZINE HYDROCHLORIDE

CONCENTRATE; ORAL

FLUPHENAZINE HYDROCHLORIDE

! PHARM ASSOC

5MG/ML

A074725 001 Sep 16, 1996

ELIXIR; ORAL

FLUPHENAZINE HYDROCHLORIDE

! PHARM ASSOC

2.5MG/5ML

A040146 001 Aug 21, 1996

INJECTABLE; INJECTION

FLUPHENAZINE HYDROCHLORIDE

! FRESENIUS KABI USA

2.5MG/ML

A089556 001 Apr 16, 1987

TABLET; ORAL

FLUPHENAZINE HYDROCHLORIDE

<u>AB</u>	AJANTA PHARMA LTD	<u>1MG</u>	<u>A217410 001</u>	Jan 05, 2023
<u>AB</u>		<u>2.5MG</u>	<u>A217410 002</u>	Jan 05, 2023
<u>AB</u>		<u>5MG</u>	<u>A217410 003</u>	Jan 05, 2023
<u>AB</u>		<u>10MG</u>	<u>A217410 004</u>	Jan 05, 2023
<u>AB</u>	ALEMBIC	<u>1MG</u>	<u>A218173 001</u>	Jul 23, 2024
<u>AB</u>		<u>2.5MG</u>	<u>A218173 002</u>	Jul 23, 2024
<u>AB</u>		<u>5MG</u>	<u>A218173 003</u>	Jul 23, 2024
<u>AB</u>		<u>10MG</u>	<u>A218173 004</u>	Jul 23, 2024
<u>AB</u>	AMNEAL	<u>1MG</u>	<u>A213647 001</u>	Jul 09, 2020
<u>AB</u>		<u>2.5MG</u>	<u>A213647 002</u>	Jul 09, 2020
<u>AB</u>		<u>5MG</u>	<u>A213647 003</u>	Jul 09, 2020
<u>AB</u>		<u>10MG</u>	<u>A213647 004</u>	Jul 09, 2020
<u>AB</u>	APOTEX	<u>1MG</u>	<u>A216649 001</u>	Jul 15, 2022
<u>AB</u>		<u>2.5MG</u>	<u>A216649 002</u>	Jul 15, 2022
<u>AB</u>		<u>5MG</u>	<u>A216649 003</u>	Jul 15, 2022
<u>AB</u>		<u>10MG</u>	<u>A216649 004</u>	Jul 15, 2022
<u>AB</u>	AUROBINDO PHARMA LTD	<u>1MG</u>	<u>A218055 001</u>	Aug 18, 2023
<u>AB</u>		<u>2.5MG</u>	<u>A218055 002</u>	Aug 18, 2023
<u>AB</u>		<u>5MG</u>	<u>A218055 003</u>	Aug 18, 2023
<u>AB</u>		<u>10MG</u>	<u>A218055 004</u>	Aug 18, 2023
<u>AB</u>	BRECKENRIDGE	<u>1MG</u>	<u>A216891 001</u>	Nov 15, 2022
<u>AB</u>		<u>2.5MG</u>	<u>A216891 002</u>	Nov 15, 2022
<u>AB</u>		<u>5MG</u>	<u>A216891 003</u>	Nov 15, 2022
<u>AB</u>		<u>10MG</u>	<u>A216891 004</u>	Nov 15, 2022
<u>AB</u>	CHARTWELL RX	<u>1MG</u>	<u>A215141 001</u>	Oct 20, 2021

PRESCRIPTION DRUG PRODUCT LIST

FLUPHENAZINE HYDROCHLORIDE

TABLET; ORAL

FLUPHENAZINE HYDROCHLORIDE

<u>AB</u>		<u>2.5MG</u>	<u>A215141 002</u>	Oct 20, 2021
<u>AB</u>		<u>5MG</u>	<u>A215141 003</u>	Oct 20, 2021
<u>AB</u>		<u>10MG</u>	<u>A215141 004</u>	Oct 20, 2021
<u>AB</u>	DR REDDYS	<u>1MG</u>	<u>A214534 001</u>	Jan 07, 2021
<u>AB</u>		<u>2.5MG</u>	<u>A214534 002</u>	Jan 07, 2021
<u>AB</u>		<u>5MG</u>	<u>A214534 003</u>	Jan 07, 2021
<u>AB</u>		<u>10MG</u>	<u>A214534 004</u>	Jan 07, 2021
<u>AB</u>	GLENMARK PHARMS LTD	<u>1MG</u>	<u>A216350 001</u>	Nov 06, 2023
<u>AB</u>		<u>2.5MG</u>	<u>A216350 002</u>	Nov 06, 2023
<u>AB</u>		<u>5MG</u>	<u>A216350 003</u>	Nov 06, 2023
<u>AB</u>		<u>10MG</u>	<u>A216350 004</u>	Nov 06, 2023
<u>AB</u>	LANNETT CO INC	<u>1MG</u>	<u>A089743 002</u>	Aug 25, 1988
<u>AB</u>		<u>2.5MG</u>	<u>A089743 003</u>	Aug 25, 1988
<u>AB</u>	!	<u>5MG</u>	<u>A089743 004</u>	Aug 25, 1988
<u>AB</u>		<u>10MG</u>	<u>A089743 001</u>	Aug 25, 1988
<u>AB</u>	MSN	<u>1MG</u>	<u>A217189 001</u>	Sep 11, 2023
<u>AB</u>		<u>2.5MG</u>	<u>A217189 002</u>	Sep 11, 2023
<u>AB</u>		<u>5MG</u>	<u>A217189 003</u>	Sep 11, 2023
<u>AB</u>		<u>10MG</u>	<u>A217189 004</u>	Sep 11, 2023
<u>AB</u>	NOVITIUM PHARMA	<u>1MG</u>	<u>A214674 001</u>	Mar 01, 2021
<u>AB</u>		<u>2.5MG</u>	<u>A214674 002</u>	Mar 01, 2021
<u>AB</u>		<u>5MG</u>	<u>A214674 003</u>	Mar 01, 2021
<u>AB</u>		<u>10MG</u>	<u>A214674 004</u>	Mar 01, 2021
<u>AB</u>	RISING	<u>1MG</u>	<u>A218283 001</u>	Jun 07, 2024
<u>AB</u>		<u>2.5MG</u>	<u>A218283 002</u>	Jun 07, 2024
<u>AB</u>		<u>5MG</u>	<u>A218283 003</u>	Jun 07, 2024
<u>AB</u>		<u>10MG</u>	<u>A218283 004</u>	Jun 07, 2024
<u>AB</u>	TARO	<u>1MG</u>	<u>A215674 001</u>	Apr 14, 2022
<u>AB</u>		<u>2.5MG</u>	<u>A215674 002</u>	Apr 14, 2022
<u>AB</u>		<u>5MG</u>	<u>A215674 003</u>	Apr 14, 2022
<u>AB</u>		<u>10MG</u>	<u>A215674 004</u>	Apr 14, 2022
<u>AB</u>	TWI PHARMS	<u>1MG</u>	<u>A215848 004</u>	Dec 14, 2022
<u>AB</u>		<u>2.5MG</u>	<u>A215848 001</u>	Apr 01, 2022
<u>AB</u>		<u>5MG</u>	<u>A215848 002</u>	Apr 01, 2022
<u>AB</u>		<u>10MG</u>	<u>A215848 003</u>	Apr 01, 2022
<u>AB</u>	UPSHER SMITH LABS	<u>1MG</u>	<u>A213784 001</u>	Oct 24, 2022
<u>AB</u>		<u>2.5MG</u>	<u>A213784 002</u>	Oct 24, 2022
<u>AB</u>		<u>5MG</u>	<u>A213784 003</u>	Oct 24, 2022
<u>AB</u>		<u>10MG</u>	<u>A213784 004</u>	Oct 24, 2022
<u>AB</u>	ZYDUS LIFESCIENCES	<u>1MG</u>	<u>A214552 001</u>	May 27, 2021
<u>AB</u>		<u>2.5MG</u>	<u>A214552 002</u>	May 27, 2021
<u>AB</u>		<u>5MG</u>	<u>A214552 003</u>	May 27, 2021
<u>AB</u>		<u>10MG</u>	<u>A214552 004</u>	May 27, 2021

FLURANDRENOLIDE

LOTION; TOPICAL

FLURANDRENOLIDE

! PADAGIS ISRAEL

0.05%

A207133 001 Aug 30, 2016

OINTMENT; TOPICAL

FLURANDRENOLIDE

! TELIGENT

0.05%

A207851 001 Dec 30, 2016

TAPE; TOPICAL

CORDRAN

+! ALMIRALL

0.004MG/SQ CM

N016455 001

FLURAZEPAM HYDROCHLORIDE

CAPSULE; ORAL

FLURAZEPAM HYDROCHLORIDE

<u>AB</u>	CHARTWELL RX	<u>15MG</u>	<u>A072368 001</u>	Mar 30, 1989
<u>AB</u>		<u>30MG</u>	<u>A072369 001</u>	Mar 30, 1989
<u>AB</u>	RISING	<u>15MG</u>	<u>A070345 002</u>	Nov 27, 1985
<u>AB</u>	!	<u>30MG</u>	<u>A070345 001</u>	Nov 27, 1985

FLURBIPROFEN

TABLET; ORAL

FLURBIPROFEN

! GENUS

100MG

A074431 001 May 31, 1995

PRESCRIPTION DRUG PRODUCT LIST

FLURBIPROFEN SODIUM

SOLUTION/DROPS;OPHTHALMIC

FLURBIPROFEN SODIUM

! BAUSCH AND LOMB

0.03%

A074447 001 Jan 04, 1995

FLURPIRIDAZ F-18

SOLUTION;INTRAVENOUS

FLYRCADO

+! GE HLTHCARE

5-55mCi/ML

N215168 001 Sep 27, 2024

FLUTAMIDE

CAPSULE;ORAL

FLUTAMIDE

! WAYLIS THERAP

125MG

A075298 001 Sep 18, 2001

FLUTEMETAMOL F-18

INJECTABLE;INTRAVENOUS

VIZAMYL

+! GE HEALTHCARE

121.5mCi/30ML (4.05mCi/ML)

N203137 002 Oct 25, 2013

FLUTICASON FUROATE

POWDER;INHALATION

ARNUITY ELLIPTA

+! GLAXOSMITHKLINE

0.05MG/INH

N205625 003 May 17, 2018

+!

0.1MG/INH

N205625 001 Aug 20, 2014

+!

0.2MG/INH

N205625 002 Aug 20, 2014

FLUTICASON FUROATE; UMECLIDINIUM BROMIDE; VILANTEROL TRIFENATATE

POWDER;INHALATION

TRELEGY ELLIPTA

+! GLAXOSMITHKLINE

0.1MG/INH;EQ 0.0625MG BASE/INH;EQ
0.025MG BASE/INH

N209482 001 Sep 18, 2017

+!

0.2MG/INH;EQ 0.0625MG BASE/INH;EQ
0.025MG BASE/INH

N209482 002 Sep 09, 2020

FLUTICASON FUROATE; VILANTEROL TRIFENATATE

POWDER;INHALATION

BREQ ELLIPTA

+ GLAXO GRP LTD

0.05MG/INH;EQ 0.025MG BASE/INH

N204275 003 May 12, 2023

+!

0.1MG/INH;EQ 0.025MG BASE/INH

N204275 001 May 10, 2013

+!

0.2MG/INH;EQ 0.025MG BASE/INH

N204275 002 Apr 30, 2015

FLUTICASON PROPIONATE

AEROSOL, METERED;INHALATION

FLOVENT HFA

+! GLAXO GRP LTD

0.044MG/INH

N021433 003 May 14, 2004

+!

0.11MG/INH

N021433 002 May 14, 2004

+!

0.22MG/INH

N021433 001 May 14, 2004

CREAM;TOPICAL

FLUTICASON PROPIONATEAB COSETTE0.05%A077055 001 Jun 30, 2006AB FOUGERA PHARMS0.05%A076451 001 May 14, 2004AB ! PADAGIS ISRAEL0.05%A076793 001 May 14, 2004

LOTION;TOPICAL

FLUTICASON PROPIONATEAB ! GLENMARK PHARMS LTD0.05%A090759 001 May 02, 2011AB PADAGIS ISRAEL0.05%A091553 001 Jul 30, 2013

OINTMENT;TOPICAL

FLUTICASON PROPIONATEAB COSETTE0.005%A077168 001 Mar 03, 2006AB ! PADAGIS ISRAEL0.005%A076668 001 May 14, 2004

POWDER;INHALATION

FLOVENT DISKUS 100

+! GLAXO GRP LTD

0.1MG/INH

N020833 002 Sep 29, 2000

FLOVENT DISKUS 250

+! GLAXO GRP LTD

0.25MG/INH

N020833 003 Sep 29, 2000

FLOVENT DISKUS 50

+! GLAXO GRP LTD

0.05MG/INH

N020833 001 Sep 29, 2000

SPRAY, METERED;NASAL

FLUTICASON PROPIONATEAB APOTEX INC0.05MG/SPRAYA077538 001 Sep 12, 2007AB CHARTWELL RX0.05MG/SPRAYA078492 001 Jan 09, 2012AB ! HIKMA0.05MG/SPRAYA076504 001 Feb 22, 2006

XHANCE

+! OPTINOSE US INC

0.093MG

N209022 001 Sep 18, 2017

PRESCRIPTION DRUG PRODUCT LIST

FLUTICASONE PROPIONATE; SALMETEROL XINAFOATE

AEROSOL, METERED; INHALATION

ADVAIR HFA

+	GLAXO GRP LTD	0.045MG/INH;EQ 0.021MG BASE/INH	N021254	001	Jun 08, 2006
+		0.115MG/INH;EQ 0.021MG BASE/INH	N021254	002	Jun 08, 2006
+		0.23MG/INH;EQ 0.021MG BASE/INH	N021254	003	Jun 08, 2006

POWDER; INHALATION

ADVAIR DISKUS 100/50

AB	+	GLAXO GRP LTD	<u>0.1MG/INH;EQ 0.05MG BASE/INH</u>	<u>N021077</u>	<u>001</u>	Aug 24, 2000
-----------	---	---------------	-------------------------------------	----------------	------------	--------------

ADVAIR DISKUS 250/50

AB	+	GLAXO GRP LTD	<u>0.25MG/INH;EQ 0.05MG BASE/INH</u>	<u>N021077</u>	<u>002</u>	Aug 24, 2000
-----------	---	---------------	--------------------------------------	----------------	------------	--------------

ADVAIR DISKUS 500/50

AB	+	GLAXO GRP LTD	<u>0.5MG/INH;EQ 0.05MG BASE/INH</u>	<u>N021077</u>	<u>003</u>	Aug 24, 2000
-----------	---	---------------	-------------------------------------	----------------	------------	--------------

FLUTICASONE PROPIONATE AND SALMETEROL XINAFOATE

AB		HIKMA	<u>0.1MG/INH;EQ 0.05MG BASE/INH</u>	<u>A203433</u>	<u>001</u>	Dec 17, 2020
-----------	--	-------	-------------------------------------	----------------	------------	--------------

AB			<u>0.25MG/INH;EQ 0.05MG BASE/INH</u>	<u>A203433</u>	<u>002</u>	Dec 17, 2020
-----------	--	--	--------------------------------------	----------------	------------	--------------

AB			<u>0.5MG/INH;EQ 0.05MG BASE/INH</u>	<u>A203433</u>	<u>003</u>	Dec 19, 2023
-----------	--	--	-------------------------------------	----------------	------------	--------------

AB		TEVA PHARMS USA	<u>0.1MG/INH;EQ 0.05MG BASE/INH</u>	<u>A213948</u>	<u>001</u>	Dec 13, 2021
-----------	--	-----------------	-------------------------------------	----------------	------------	--------------

AB			<u>0.25MG/INH;EQ 0.05MG BASE/INH</u>	<u>A213948</u>	<u>002</u>	Dec 13, 2021
-----------	--	--	--------------------------------------	----------------	------------	--------------

AB			<u>0.5MG/INH;EQ 0.05MG BASE/INH</u>	<u>A213948</u>	<u>003</u>	Dec 13, 2021
-----------	--	--	-------------------------------------	----------------	------------	--------------

WIXELA INHUB

AB		MYLAN	<u>0.1MG/INH;EQ 0.05MG BASE/INH</u>	<u>A208891</u>	<u>001</u>	Jan 30, 2019
-----------	--	-------	-------------------------------------	----------------	------------	--------------

AB			<u>0.25MG/INH;EQ 0.05MG BASE/INH</u>	<u>A208891</u>	<u>002</u>	Jan 30, 2019
-----------	--	--	--------------------------------------	----------------	------------	--------------

AB			<u>0.5MG/INH;EQ 0.05MG BASE/INH</u>	<u>A208891</u>	<u>003</u>	Jan 30, 2019
-----------	--	--	-------------------------------------	----------------	------------	--------------

AIRDUO RESPICLICK

+	TEVA PHARM	0.055MG/INH;EQ 0.014MG BASE/INH	N208799	001	Jan 27, 2017
---	------------	---------------------------------	---------	-----	--------------

+		0.113MG/INH;EQ 0.014MG BASE/INH	N208799	002	Jan 27, 2017
---	--	---------------------------------	---------	-----	--------------

+		0.232MG/INH;EQ 0.014MG BASE/INH	N208799	003	Jan 27, 2017
---	--	---------------------------------	---------	-----	--------------

FLUVASTATIN SODIUM

CAPSULE; ORAL

FLUVASTATIN SODIUM

AB		MYLAN PHARMS INC	<u>EQ 20MG BASE</u>	<u>A090595</u>	<u>001</u>	Apr 11, 2012
-----------	--	------------------	---------------------	----------------	------------	--------------

AB	!		<u>EQ 40MG BASE</u>	<u>A090595</u>	<u>002</u>	Apr 11, 2012
-----------	---	--	---------------------	----------------	------------	--------------

AB		TEVA PHARMS	<u>EQ 20MG BASE</u>	<u>A078407</u>	<u>001</u>	Jun 12, 2012
-----------	--	-------------	---------------------	----------------	------------	--------------

AB			<u>EQ 40MG BASE</u>	<u>A078407</u>	<u>002</u>	Jun 12, 2012
-----------	--	--	---------------------	----------------	------------	--------------

TABLET, EXTENDED RELEASE; ORAL

FLUVASTATIN SODIUM

AB		BEIJING	<u>EQ 80MG BASE</u>	<u>A209397</u>	<u>001</u>	Apr 26, 2021
-----------	--	---------	---------------------	----------------	------------	--------------

AB		TEVA PHARMS USA	<u>EQ 80MG BASE</u>	<u>A079011</u>	<u>001</u>	Jan 27, 2016
-----------	--	-----------------	---------------------	----------------	------------	--------------

LESCOL XL

AB	+	SANDOZ	<u>EQ 80MG BASE</u>	<u>N021192</u>	<u>001</u>	Oct 06, 2000
-----------	---	--------	---------------------	----------------	------------	--------------

FLUVOXAMINE MALEATE

CAPSULE, EXTENDED RELEASE; ORAL

FLUVOXAMINE MALEATE

AB		ACTAVIS ELIZABETH	<u>100MG</u>	<u>A091482</u>	<u>001</u>	Apr 23, 2013
-----------	--	-------------------	--------------	----------------	------------	--------------

AB	!		<u>150MG</u>	<u>A091482</u>	<u>002</u>	Nov 18, 2013
-----------	---	--	--------------	----------------	------------	--------------

AB		AJANTA PHARMA LTD	<u>100MG</u>	<u>A219055</u>	<u>001</u>	Oct 17, 2024
-----------	--	-------------------	--------------	----------------	------------	--------------

AB			<u>150MG</u>	<u>A219055</u>	<u>002</u>	Oct 17, 2024
-----------	--	--	--------------	----------------	------------	--------------

AB		BIONPHARMA	<u>100MG</u>	<u>A212182</u>	<u>002</u>	Sep 16, 2020
-----------	--	------------	--------------	----------------	------------	--------------

AB			<u>150MG</u>	<u>A212182</u>	<u>001</u>	May 11, 2020
-----------	--	--	--------------	----------------	------------	--------------

AB		ENDO OPERATIONS	<u>100MG</u>	<u>A091476</u>	<u>001</u>	Mar 13, 2013
-----------	--	-----------------	--------------	----------------	------------	--------------

AB			<u>150MG</u>	<u>A091476</u>	<u>002</u>	Mar 13, 2013
-----------	--	--	--------------	----------------	------------	--------------

TABLET; ORAL

FLUVOXAMINE MALEATE

AB		APOTEX	<u>25MG</u>	<u>A075902</u>	<u>001</u>	May 07, 2001
-----------	--	--------	-------------	----------------	------------	--------------

AB			<u>50MG</u>	<u>A075902</u>	<u>002</u>	May 07, 2001
-----------	--	--	-------------	----------------	------------	--------------

AB			<u>100MG</u>	<u>A075902</u>	<u>003</u>	May 07, 2001
-----------	--	--	--------------	----------------	------------	--------------

AB		BIONPHARMA	<u>25MG</u>	<u>A217917</u>	<u>001</u>	Jan 22, 2024
-----------	--	------------	-------------	----------------	------------	--------------

AB			<u>50MG</u>	<u>A217917</u>	<u>002</u>	Jan 22, 2024
-----------	--	--	-------------	----------------	------------	--------------

AB			<u>100MG</u>	<u>A217917</u>	<u>003</u>	Jan 22, 2024
-----------	--	--	--------------	----------------	------------	--------------

AB		UPSHER SMITH LABS	<u>25MG</u>	<u>A075888</u>	<u>001</u>	Nov 29, 2000
-----------	--	-------------------	-------------	----------------	------------	--------------

AB			<u>50MG</u>	<u>A075888</u>	<u>002</u>	Nov 29, 2000
-----------	--	--	-------------	----------------	------------	--------------

AB	!		<u>100MG</u>	<u>A075888</u>	<u>003</u>	Nov 29, 2000
-----------	---	--	--------------	----------------	------------	--------------

LUVOX

AB	+	ANI PHARMS	<u>25MG</u>	<u>N021519</u>	<u>001</u>	Dec 20, 2007
-----------	---	------------	-------------	----------------	------------	--------------

AB	+		<u>50MG</u>	<u>N021519</u>	<u>002</u>	Dec 20, 2007
-----------	---	--	-------------	----------------	------------	--------------

AB	+		<u>100MG</u>	<u>N021519</u>	<u>003</u>	Dec 20, 2007
-----------	---	--	--------------	----------------	------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

FOLIC ACID

INJECTABLE; INJECTION

FOLIC ACID

<u>AP</u>	!	FRESENIUS KABI USA	<u>5MG/ML</u>	<u>A089202</u>	<u>001</u>	Feb 18, 1986
<u>AP</u>		XGEN PHARMS	<u>5MG/ML</u>	<u>A202522</u>	<u>001</u>	Nov 06, 2019

TABLET; ORAL

FOLIC ACID

<u>AA</u>	!	AMNEAL PHARM	<u>1MG</u>	<u>A040625</u>	<u>001</u>	Jul 21, 2005
<u>AA</u>		ATHEM	<u>1MG</u>	<u>A211064</u>	<u>001</u>	Mar 08, 2019
<u>AA</u>		CADILA PHARMS LTD	<u>1MG</u>	<u>A202437</u>	<u>001</u>	Jan 27, 2014
<u>AA</u>		CHARTWELL MOLECULAR	<u>1MG</u>	<u>A090035</u>	<u>001</u>	Jun 09, 2009
<u>AA</u>		LEADING	<u>1MG</u>	<u>A040796</u>	<u>001</u>	Jan 12, 2009
<u>AA</u>		NUVO PHARMS INC	<u>1MG</u>	<u>A204418</u>	<u>001</u>	Jul 28, 2015
<u>AA</u>		QINGDAO BAHEAL PHARM	<u>1MG</u>	<u>A091145</u>	<u>001</u>	Jul 12, 2013
<u>AA</u>	+	WATSON LABS	<u>1MG</u>	<u>A080680</u>	<u>001</u>	

FOMEPIZOLE

INJECTABLE; INJECTION

FOMEPIZOLE

<u>AP</u>		AM REGENT	<u>1.5GM/1.5ML (1GM/ML)</u>	<u>A078368</u>	<u>001</u>	Dec 14, 2007
<u>AP</u>		GLAND PHARMA LTD	<u>1.5GM/1.5ML (1GM/ML)</u>	<u>A216791</u>	<u>001</u>	Jul 06, 2023
<u>AP</u>	!	MYLAN INSTITUTIONAL	<u>1.5GM/1.5ML (1GM/ML)</u>	<u>A078639</u>	<u>001</u>	Mar 03, 2008
<u>AP</u>		NAVINTA LLC	<u>1.5GM/1.5ML (1GM/ML)</u>	<u>A078537</u>	<u>001</u>	Mar 06, 2008

FONDAPARINUX SODIUM

INJECTABLE; SUBCUTANEOUS

ARIXTRA

<u>AP</u>	+	MYLAN IRELAND LTD	<u>2.5MG/0.5ML</u>	<u>N021345</u>	<u>001</u>	Dec 07, 2001
<u>AP</u>	+		<u>5MG/0.4ML</u>	<u>N021345</u>	<u>002</u>	May 28, 2004
<u>AP</u>	+		<u>7.5MG/0.6ML</u>	<u>N021345</u>	<u>003</u>	May 28, 2004
<u>AP</u>	+		<u>10MG/0.8ML</u>	<u>N021345</u>	<u>004</u>	May 28, 2004

FONDAPARINUX SODIUM

<u>AP</u>		BRIGHTGENE	<u>2.5MG/0.5ML</u>	<u>A218312</u>	<u>001</u>	Dec 18, 2024
<u>AP</u>		DR REDDYS LABS LTD	<u>2.5MG/0.5ML</u>	<u>A091316</u>	<u>001</u>	Jul 11, 2011
<u>AP</u>			<u>5MG/0.4ML</u>	<u>A091316</u>	<u>002</u>	Jul 11, 2011
<u>AP</u>			<u>7.5MG/0.6ML</u>	<u>A091316</u>	<u>003</u>	Jul 11, 2011
<u>AP</u>			<u>10MG/0.8ML</u>	<u>A091316</u>	<u>004</u>	Jul 11, 2011
<u>AP</u>		EUGIA PHARMA	<u>2.5MG/0.5ML</u>	<u>A206918</u>	<u>001</u>	Dec 26, 2017
<u>AP</u>			<u>5MG/0.4ML</u>	<u>A206918</u>	<u>002</u>	Dec 26, 2017
<u>AP</u>			<u>7.5MG/0.6ML</u>	<u>A206918</u>	<u>003</u>	Dec 26, 2017
<u>AP</u>			<u>10MG/0.8ML</u>	<u>A206918</u>	<u>004</u>	Dec 26, 2017
<u>AP</u>		HANGZHOU ZHONGMEI	<u>2.5MG/0.5ML</u>	<u>A216493</u>	<u>001</u>	Aug 19, 2024
<u>AP</u>			<u>5MG/0.4ML</u>	<u>A216493</u>	<u>002</u>	Aug 19, 2024
<u>AP</u>			<u>7.5MG/0.6ML</u>	<u>A216493</u>	<u>003</u>	Aug 19, 2024
<u>AP</u>			<u>10MG/0.8ML</u>	<u>A216493</u>	<u>004</u>	Aug 19, 2024
<u>AP</u>		HENGRUI PHARMA	<u>2.5MG/0.5ML</u>	<u>A206812</u>	<u>001</u>	May 15, 2018
<u>AP</u>			<u>5MG/0.4ML</u>	<u>A206812</u>	<u>002</u>	May 15, 2018
<u>AP</u>			<u>7.5MG/0.6ML</u>	<u>A206812</u>	<u>003</u>	May 15, 2018
<u>AP</u>			<u>10MG/0.8ML</u>	<u>A206812</u>	<u>004</u>	May 15, 2018
<u>AP</u>		SCINOPHARM TAIWAN	<u>2.5MG/0.5ML</u>	<u>A208615</u>	<u>001</u>	Nov 14, 2018
<u>AP</u>			<u>5MG/0.4ML</u>	<u>A208615</u>	<u>002</u>	Nov 14, 2018
<u>AP</u>			<u>7.5MG/0.6ML</u>	<u>A208615</u>	<u>003</u>	Nov 14, 2018
<u>AP</u>			<u>10MG/0.8ML</u>	<u>A208615</u>	<u>004</u>	Nov 14, 2018

FORMOTEROL FUMARATE

SOLUTION; INHALATION

FORMOTEROL FUMARATE

<u>AN</u>		ALEMBIC	<u>0.02MG/2ML</u>	<u>A215078</u>	<u>001</u>	Nov 22, 2021
<u>AN</u>		LEXENPHARM	<u>0.02MG/2ML</u>	<u>A216426</u>	<u>001</u>	Oct 22, 2024
<u>AN</u>		LUPIN	<u>0.02MG/2ML</u>	<u>A215053</u>	<u>001</u>	Aug 22, 2022
<u>AN</u>		MANKIND PHARMA	<u>0.02MG/2ML</u>	<u>A215883</u>	<u>001</u>	Mar 22, 2023
<u>AN</u>		MICRO LABS	<u>0.02MG/2ML</u>	<u>A218304</u>	<u>001</u>	Jun 26, 2024
<u>AN</u>		RITEDOSE CORP	<u>0.02MG/2ML</u>	<u>A216486</u>	<u>001</u>	Nov 25, 2022
<u>AN</u>		TEVA PHARMS USA INC	<u>0.02MG/2ML</u>	<u>A091141</u>	<u>001</u>	Jun 22, 2021
<u>AN</u>		WILSHIRE PHARMS INC	<u>0.02MG/2ML</u>	<u>A215621</u>	<u>001</u>	Dec 13, 2022
<u>AN</u>	+	MYLAN SPECLT	<u>0.02MG/2ML</u>	<u>N022007</u>	<u>001</u>	May 11, 2007

PRESCRIPTION DRUG PRODUCT LIST

FORMOTEROL FUMARATE; GLYCOPYRROLATE

AEROSOL, METERED; INHALATION

BEVESPI AEROSPHERE

+! ASTRAZENECA 0.0048MG/INH;0.0090MG/INH N208294 001 Apr 25, 2016

FORMOTEROL FUMARATE; MOMETASONE FUROATE

AEROSOL, METERED; INHALATION

DULERA

+! ORGANON LLC 0.005MG/INH;0.05MG/INH N022518 003 Aug 12, 2019

+! 0.005MG/INH;0.1MG/INH N022518 001 Jun 22, 2010

+! 0.005MG/INH;0.2MG/INH N022518 002 Jun 22, 2010

FOSAMPRENAVIR CALCIUM

TABLET; ORAL

FOSAMPRENAVIR CALCIUM**AB** ! MYLAN **EQ 700MG BASE** **A204060 001** Apr 15, 2016**AB** SUN PHARM **EQ 700MG BASE** **A204024 001** Nov 20, 2019FOSAPREPITANT DIMEGLUMINE

POWDER; INTRAVENOUS

EMEND**AP** +! MERCK AND CO INC **EQ 150MG BASE/VIAL** **N022023 002** Nov 12, 2010FOSAPREPITANT DIMEGLUMINE**AP** ASPIRO **EQ 150MG BASE/VIAL** **A214616 001** Jul 29, 2021**AP** BAXTER HLTHCARE **EQ 150MG BASE/VIAL** **A211860 001** Sep 05, 2019

CORP

AP BE PHARMS **EQ 150MG BASE/VIAL** **A212309 001** Sep 05, 2019**AP** CHIA TAI TIANQING **EQ 150MG BASE/VIAL** **A212143 001** Mar 03, 2021**AP** DR REDDYS **EQ 150MG BASE/VIAL** **A211160 001** Dec 09, 2020**AP** EUGIA PHARMA **EQ 150MG BASE/VIAL** **A210625 001** Jan 12, 2021**AP** FRESENIUS KABI USA **EQ 150MG BASE/VIAL** **A206197 001** Jun 09, 2016**AP** LUPIN LTD **EQ 150MG BASE/VIAL** **A210689 001** Sep 05, 2019**AP** MSN **EQ 150MG BASE/VIAL** **A209965 001** Sep 05, 2019**AP** MYLAN LABS LTD **EQ 150MG BASE/VIAL** **A204015 002** Sep 05, 2019**AP** PIRAMAL CRITICAL **EQ 150MG BASE/VIAL** **A214683 001** May 16, 2023**AP** QILU PHARM HAINAN **EQ 150MG BASE/VIAL** **A213106 001** Sep 08, 2020

TEVA PHARMS USA EQ 150MG BASE/VIAL N210064 001 Sep 05, 2019

SOLUTION; INTRAVENOUS

FOCINVEZ

+! STERISCIENCE EQ 150MG BASE/50ML (EQ 3MG BASE/ML) N216686 001 Aug 22, 2023

FOSCARBIDOPA; FOSLEVODOPA

SOLUTION; SUBCUTANEOUS

VYALEV

+! ABBVIE 120MG/10ML;2400MG/10ML N216962 001 Oct 16, 2024

(12MG/ML;240MG/ML)

FOSCARNET SODIUM

INJECTABLE; INJECTION

FOSCARNET SODIUM**AP** AMNEAL **2.4GM/100ML** **A216602 001** Mar 01, 2024**AP** AVET LIFESCIENCES **2.4GM/100ML** **A213807 001** Jun 05, 2023**AP** FRESENIUS KABI USA **2.4GM/100ML** **A212483 001** Jan 29, 2021**AP** GLAND PHARMA LTD **2.4GM/100ML** **A213001 001** Apr 21, 2021**AP** SCIECURE **2.4GM/100ML** **A213987 001** Nov 29, 2023FOSCAVIR**AP** +! CLINIGEN HLTHCARE **2.4GM/100ML** **N020068 001** Sep 27, 1991FOSDENOPTERIN HYDROBROMIDE

POWDER; INTRAVENOUS

NULIBRY

+! SENTYNL THERAPS INC EQ 9.5MG BASE/VIAL N214018 001 Feb 26, 2021

FOSFOMYCIN TROMETHAMINE

FOR SOLUTION; ORAL

FOSFOMYCIN TROMETHAMINE**AA** ALKEM LABS LTD **EQ 3GM BASE/PACKET** **A214554 001** Oct 21, 2021**AA** AMNEAL **EQ 3GM BASE/PACKET** **A216600 001** Mar 25, 2024**AA** CIPLA **EQ 3GM BASE/PACKET** **A211881 001** Jan 26, 2022**AA** ! XIROMED **EQ 3GM BASE/PACKET** **A212548 001** Oct 06, 2020

PRESCRIPTION DRUG PRODUCT LIST

FOSINOPRIL SODIUM

TABLET; ORAL

FOSINOPRIL SODIUM

<u>AB</u>	APOTEX INC	<u>10MG</u>	<u>A076906 001</u>	May 17, 2005
<u>AB</u>		<u>20MG</u>	<u>A076906 002</u>	May 17, 2005
<u>AB</u>		<u>40MG</u>	<u>A076906 003</u>	May 17, 2005
<u>AB</u>	AUROBINDO PHARMA LTD	<u>10MG</u>	<u>A091163 001</u>	Mar 30, 2011
<u>AB</u>		<u>20MG</u>	<u>A091163 002</u>	Mar 30, 2011
<u>AB</u>		<u>40MG</u>	<u>A091163 003</u>	Mar 30, 2011
<u>AB</u>	CHARTWELL RX	<u>10MG</u>	<u>A076483 001</u>	Apr 23, 2004
<u>AB</u>		<u>20MG</u>	<u>A076483 002</u>	Apr 23, 2004
<u>AB</u>		<u>40MG</u>	<u>A076483 003</u>	Apr 23, 2004
<u>AB</u>	INVAGEN PHARMS	<u>10MG</u>	<u>A077222 001</u>	Apr 20, 2005
<u>AB</u>		<u>20MG</u>	<u>A077222 002</u>	Apr 20, 2005
<u>AB</u>		<u>40MG</u>	<u>A077222 003</u>	Apr 20, 2005
<u>AB</u>	PRINSTON INC	<u>10MG</u>	<u>A205670 001</u>	Aug 29, 2016
<u>AB</u>		<u>20MG</u>	<u>A205670 002</u>	Aug 29, 2016
<u>AB</u>		<u>40MG</u>	<u>A205670 003</u>	Aug 29, 2016
<u>AB</u>	TEVA	<u>10MG</u>	<u>A076139 001</u>	Nov 25, 2003
<u>AB</u>		<u>20MG</u>	<u>A076139 002</u>	Nov 25, 2003
<u>AB</u>	!	<u>40MG</u>	<u>A076139 003</u>	Nov 25, 2003

FOSINOPRIL SODIUM; HYDROCHLOROTHIAZIDE

TABLET; ORAL

FOSINOPRIL SODIUM AND HYDROCHLOROTHIAZIDE

<u>AB</u>	AUROBINDO PHARMA	<u>10MG;12.5MG</u>	<u>A079245 001</u>	Jul 09, 2009
<u>AB</u>	!	<u>20MG;12.5MG</u>	<u>A079245 002</u>	Jul 09, 2009
<u>AB</u>	INVAGEN PHARMS	<u>10MG;12.5MG</u>	<u>A090228 001</u>	Jul 09, 2009
<u>AB</u>		<u>20MG;12.5MG</u>	<u>A090228 002</u>	Jul 09, 2009
<u>AB</u>	SANDOZ	<u>10MG;12.5MG</u>	<u>A076961 001</u>	Sep 28, 2005
<u>AB</u>		<u>20MG;12.5MG</u>	<u>A076961 002</u>	Sep 28, 2005

FOSNETUPITANT CHLORIDE HYDROCHLORIDE; PALONOSETRON HYDROCHLORIDE

SOLUTION; INTRAVENOUS

AKYNZEO

+	!	HELSINN HLTHCARE	EQ 235MG BASE/20ML (EQ 11.75MG BASE/ML);EQ 0.25MG BASE/20ML (EQ 0.0125MG BASE/ML)	N210493 002	May 27, 2020
---	---	------------------	---	-------------	--------------

FOSPHENYTOIN SODIUM

INJECTABLE; INJECTION

CEREBYX

<u>AP</u>	+	!	PARKE DAVIS	<u>EQ 50MG PHENYTOIN NA/ML</u>	<u>N020450 001</u>	Aug 05, 1996
<u>FOSPHENYTOIN SODIUM</u>						
<u>AP</u>			FRESENIUS KABI USA	<u>EQ 50MG PHENYTOIN NA/ML</u>	<u>A078052 001</u>	Aug 06, 2007
<u>AP</u>			GLAND PHARMA LTD	<u>EQ 50MG PHENYTOIN NA/ML</u>	<u>A214926 001</u>	Oct 13, 2023
<u>AP</u>			HIKMA	<u>EQ 50MG PHENYTOIN NA/ML</u>	<u>A077481 001</u>	Aug 06, 2007
<u>AP</u>				<u>EQ 50MG PHENYTOIN NA/ML</u>	<u>A077989 001</u>	Aug 06, 2007
<u>AP</u>			HIKMA FARMACEUTICA	<u>EQ 50MG PHENYTOIN NA/ML</u>	<u>A078765 001</u>	Dec 02, 2009
<u>AP</u>			SUN PHARM	<u>EQ 50MG PHENYTOIN NA/ML</u>	<u>A078417 001</u>	Mar 18, 2008
<u>AP</u>			WOCKHARDT	<u>EQ 50MG PHENYTOIN NA/ML</u>	<u>A078137 001</u>	Aug 06, 2007

FOSTAMATINIB DISODIUM

TABLET; ORAL

TAVALISSE

+		RIGEL PHARMS	EQ 100MG BASE	N209299 001	Apr 17, 2018
+	!		EQ 150MG BASE	N209299 002	Apr 17, 2018

FOSTEMSAVIR TROMETHAMINE

TABLET, EXTENDED RELEASE; ORAL

RUKOBIA

+	!	VIIV HLTHCARE	EQ 600MG BASE	N212950 001	Jul 02, 2020
---	---	---------------	---------------	-------------	--------------

FROVATRIPTAN SUCCINATE

TABLET; ORAL

FROVA

<u>AB</u>	+	!	ENDO OPERATIONS	<u>EQ 2.5MG BASE</u>	<u>N021006 001</u>	Nov 08, 2001
<u>FROVATRIPTAN SUCCINATE</u>						
<u>AB</u>			AMNEAL PHARMS CO	<u>EQ 2.5MG BASE</u>	<u>A211292 001</u>	Nov 06, 2018
<u>AB</u>			GLENMARK PHARMS LTD	<u>EQ 2.5MG BASE</u>	<u>A204730 001</u>	Mar 11, 2016
<u>AB</u>			RENATA	<u>EQ 2.5MG BASE</u>	<u>A213891 001</u>	Apr 06, 2022

PRESCRIPTION DRUG PRODUCT LIST

FRUQUINTINIB

CAPSULE; ORAL

FRUZAQLA

+ TAKEDA PHARMS USA

1MG

N217564 001 Nov 08, 2023

+!

5MG

N217564 002 Nov 08, 2023

FULVESTRANT

SOLUTION; INTRAMUSCULAR

FASLODEXAO +! ASTRAZENECA250MG/5ML (50MG/ML)N021344 001 Apr 25, 2002FULVESTRANTAO ACCORD HLTHCARE250MG/5ML (50MG/ML)A211689 001 Nov 17, 2020AO ALEMBIC250MG/5ML (50MG/ML)A215077 001 Dec 22, 2022AO AMNEAL250MG/5ML (50MG/ML)A210044 001 Mar 04, 2019AO CHIA TAI TIANQING250MG/5ML (50MG/ML)A211422 001 Feb 07, 2020AO DR REDDYS250MG/5ML (50MG/ML)A209246 001 Aug 07, 2020AO EUGIA PHARMA250MG/5ML (50MG/ML)A208811 001 Jul 23, 2019AO250MG/5ML (50MG/ML)A215169 001 Jun 30, 2023AO FRESENIUS KABI USA250MG/5ML (50MG/ML)N210326 001 May 20, 2019AO GLENMARK PHARMS INC250MG/5ML (50MG/ML)A207754 001 Aug 22, 2019AO HBT LABS INC250MG/5ML (50MG/ML)A209714 001 Nov 21, 2019AO JIANGSU HANSOH250MG/5ML (50MG/ML)A214682 001 Feb 10, 2022

PHARM

AO SAGENT PHARMS INC250MG/5ML (50MG/ML)A205871 001 Aug 22, 2019AO SANDOZ250MG/5ML (50MG/ML)A205935 001 May 14, 2019AO XIROMED250MG/5ML (50MG/ML)A213553 001 Aug 13, 2021AO ZYDUS PHARMS250MG/5ML (50MG/ML)A215234 001 Jul 29, 2021FUROSEMIDE

INJECTABLE; INJECTION

FUROSEMIDEAP ACCORD HLTHCARE10MG/MLA070017 001 Dec 15, 1986AP AMNEAL PHARMS CO10MG/MLA207552 001 Jul 20, 2016AP AREVA PHARMS10MG/MLA208435 001 Dec 18, 2020AP ASPIRO10MG/MLA217557 001 Dec 13, 2024AP AVET LIFESCIENCES10MG/MLA203428 001 Aug 26, 2014AP ! BAXTER HLTHCARE10MG/MLA202747 001 Jan 27, 2014

CORP

AP EUGIA PHARMA10MG/MLA212174 001 May 03, 2019AP FRESENIUS KABI USA10MG/MLN018902 001 May 22, 1984AP GLAND PHARMA LTD10MG/MLA213902 001 Jul 01, 2020AP HIKMA10MG/MLN018267 001AP HOSPIRA10MG/MLA075241 001 May 28, 1999AP10MG/MLN018667 001 May 28, 1982AP MANKIND PHARMA10MG/MLA216860 001 Dec 16, 2022AP MEITHEAL10MG/MLA212803 001 Jan 27, 2022AP SABA ILAC SANAYIVE10MG/MLA215856 001 Apr 02, 2024AP SAGENT10MG/MLA214766 001 Jan 27, 2021AP WOCKHARDT10MG/MLA077941 001 Mar 22, 2007

SOLUTION; ORAL

FUROSEMIDE

! HIKMA

10MG/ML

A070434 001 Apr 22, 1987

40MG/5ML

A070433 001 Apr 22, 1987

SOLUTION; SUBCUTANEOUS

FUROSCIX

+! SCPHARMACEUTICALS

80MG/10ML (8MG/ML)

N209988 001 Oct 07, 2022

TABLET; ORAL

FUROSEMIDEAB EPIC PHARMA LLC20MGN018569 002AB40MGN018569 001AB80MGN018569 005 Aug 14, 1984AB GRAVITI PHARMS20MGA216629 001 Oct 17, 2022AB40MGA216629 002 Oct 17, 2022AB80MGA216629 003 Oct 17, 2022AB HIKMA20MGN018823 001 Nov 10, 1983AB40MGN018823 002 Nov 10, 1983AB80MGA070086 001 Jan 24, 1986AB IPCA LABS LTD20MGA078010 001 Sep 18, 2006AB40MGA078010 002 Sep 18, 2006AB80MGA078010 003 Sep 18, 2006AB LEADING20MGA077293 001 Nov 09, 2005AB40MGA077293 002 Nov 09, 2005AB80MGA077293 003 Nov 09, 2005AB PRINSTON INC20MGA076796 001 Mar 26, 2004

PRESCRIPTION DRUG PRODUCT LIST

FUROSEMIDE

TABLET; ORAL

FUROSEMIDE

<u>AB</u>		<u>40MG</u>	<u>A076796</u>	<u>002</u>	Mar 26, 2004
<u>AB</u>		<u>80MG</u>	<u>A076796</u>	<u>003</u>	Mar 26, 2004

LASIX

<u>AB</u>	+	VALIDUS PHARMS	<u>20MG</u>	<u>N016273</u>	<u>002</u>
<u>AB</u>	+		<u>40MG</u>	<u>N016273</u>	<u>001</u>
<u>AB</u>	+	!	<u>80MG</u>	<u>N016273</u>	<u>003</u>

FUTIBATINIB

TABLET; ORAL

LYTGOBI

+	!	TAIHO ONCOLOGY	4MG	N214801	001	Sep 30, 2022
---	---	----------------	-----	---------	-----	--------------

GABAPENTIN

CAPSULE; ORAL

GABAPENTIN

<u>AB</u>		ACTAVIS ELIZABETH	<u>100MG</u>	<u>A075350</u>	<u>001</u>	Sep 12, 2003
<u>AB</u>			<u>300MG</u>	<u>A075350</u>	<u>002</u>	Sep 12, 2003
<u>AB</u>			<u>400MG</u>	<u>A075350</u>	<u>003</u>	Sep 12, 2003
<u>AB</u>		ALKEM	<u>100MG</u>	<u>A090858</u>	<u>001</u>	Dec 17, 2010
<u>AB</u>			<u>300MG</u>	<u>A090858</u>	<u>002</u>	Dec 17, 2010
<u>AB</u>			<u>400MG</u>	<u>A090858</u>	<u>003</u>	Dec 17, 2010
<u>AB</u>		AMNEAL PHARMS NY	<u>100MG</u>	<u>A078428</u>	<u>001</u>	Jul 25, 2007
<u>AB</u>			<u>300MG</u>	<u>A078428</u>	<u>002</u>	Jul 25, 2007
<u>AB</u>			<u>400MG</u>	<u>A078428</u>	<u>003</u>	Jul 25, 2007
<u>AB</u>		ASCENT PHARMS INC	<u>100MG</u>	<u>A214956</u>	<u>001</u>	May 10, 2021
<u>AB</u>			<u>300MG</u>	<u>A214956</u>	<u>002</u>	May 10, 2021
<u>AB</u>			<u>400MG</u>	<u>A214956</u>	<u>003</u>	May 10, 2021
<u>AB</u>		AUROBINDO PHARMA	<u>100MG</u>	<u>A078787</u>	<u>001</u>	Jan 31, 2008
<u>AB</u>			<u>300MG</u>	<u>A078787</u>	<u>002</u>	Jan 31, 2008
<u>AB</u>			<u>400MG</u>	<u>A078787</u>	<u>003</u>	Jan 31, 2008
<u>AB</u>		CSPC OUYI	<u>100MG</u>	<u>A075477</u>	<u>001</u>	Mar 23, 2005
<u>AB</u>			<u>300MG</u>	<u>A075477</u>	<u>002</u>	Mar 23, 2005
<u>AB</u>			<u>400MG</u>	<u>A075477</u>	<u>003</u>	Mar 23, 2005
<u>AB</u>		GRANULES	<u>100MG</u>	<u>A075360</u>	<u>001</u>	Apr 06, 2005
<u>AB</u>			<u>300MG</u>	<u>A075360</u>	<u>002</u>	Apr 06, 2005
<u>AB</u>			<u>400MG</u>	<u>A075360</u>	<u>003</u>	Apr 06, 2005
<u>AB</u>		INVAGEN PHARMS	<u>100MG</u>	<u>A090705</u>	<u>001</u>	Dec 30, 2009
<u>AB</u>			<u>300MG</u>	<u>A090705</u>	<u>002</u>	Dec 30, 2009
<u>AB</u>			<u>400MG</u>	<u>A090705</u>	<u>003</u>	Dec 30, 2009
<u>AB</u>		LAURUS	<u>100MG</u>	<u>A217546</u>	<u>001</u>	May 12, 2023
<u>AB</u>			<u>300MG</u>	<u>A217546</u>	<u>002</u>	May 12, 2023
<u>AB</u>			<u>400MG</u>	<u>A217546</u>	<u>003</u>	May 12, 2023
<u>AB</u>		MARKSANS PHARMA	<u>100MG</u>	<u>A090007</u>	<u>001</u>	Jul 21, 2011
<u>AB</u>			<u>300MG</u>	<u>A090007</u>	<u>002</u>	Jul 21, 2011
<u>AB</u>			<u>400MG</u>	<u>A090007</u>	<u>003</u>	Jul 21, 2011
<u>AB</u>		RISING	<u>100MG</u>	<u>A207099</u>	<u>001</u>	Mar 24, 2017
<u>AB</u>			<u>300MG</u>	<u>A207099</u>	<u>002</u>	Mar 24, 2017
<u>AB</u>			<u>400MG</u>	<u>A207099</u>	<u>003</u>	Mar 24, 2017
<u>AB</u>		SCIEGEN PHARMS INC	<u>100MG</u>	<u>A204989</u>	<u>001</u>	Feb 18, 2016
<u>AB</u>			<u>300MG</u>	<u>A204989</u>	<u>002</u>	Feb 18, 2016
<u>AB</u>			<u>400MG</u>	<u>A204989</u>	<u>003</u>	Feb 18, 2016
<u>AB</u>		STRIDES PHARMA	<u>100MG</u>	<u>A206943</u>	<u>001</u>	May 14, 2018
<u>AB</u>			<u>100MG</u>	<u>A211314</u>	<u>001</u>	Oct 16, 2018
<u>AB</u>			<u>300MG</u>	<u>A206943</u>	<u>002</u>	May 14, 2018
<u>AB</u>			<u>300MG</u>	<u>A211314</u>	<u>002</u>	Oct 16, 2018
<u>AB</u>			<u>400MG</u>	<u>A206943</u>	<u>003</u>	May 14, 2018
<u>AB</u>			<u>400MG</u>	<u>A211314</u>	<u>003</u>	Oct 16, 2018
<u>AB</u>		TARO	<u>100MG</u>	<u>A077261</u>	<u>001</u>	Aug 02, 2013
<u>AB</u>			<u>300MG</u>	<u>A077261</u>	<u>002</u>	Aug 02, 2013
<u>AB</u>			<u>400MG</u>	<u>A077261</u>	<u>003</u>	Aug 02, 2013
<u>AB</u>		ZHEJIANG YONGTAI	<u>100MG</u>	<u>A213603</u>	<u>001</u>	Aug 17, 2020
<u>AB</u>			<u>300MG</u>	<u>A213603</u>	<u>002</u>	Aug 17, 2020
<u>AB</u>			<u>400MG</u>	<u>A213603</u>	<u>003</u>	Aug 17, 2020

NEURONTIN

<u>AB</u>	+	VIATRIS	<u>100MG</u>	<u>N020235</u>	<u>001</u>	Dec 30, 1993
<u>AB</u>	+		<u>300MG</u>	<u>N020235</u>	<u>002</u>	Dec 30, 1993
<u>AB</u>	+	!	<u>400MG</u>	<u>N020235</u>	<u>003</u>	Dec 30, 1993

SOLUTION; ORAL

GABAPENTIN

<u>AA</u>		ACELLA PHARMS LLC	<u>250MG/5ML</u>	<u>A076403</u>	<u>001</u>	May 01, 2012
-----------	--	-------------------	------------------	----------------	------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

GABAPENTIN

SOLUTION;ORAL

GABAPENTIN

<u>AA</u>	AMNEAL PHARMS	<u>250MG/5ML</u>	<u>A202024 001</u>	Mar 23, 2012
<u>AA</u>	ANNORA PHARMA	<u>250MG/5ML</u>	<u>A217682 001</u>	Jan 17, 2024
<u>AA</u>	BELCHER	<u>250MG/5ML</u>	<u>A091286 001</u>	Mar 14, 2016
<u>AA</u>	MISSION PHARMACAL	<u>250MG/5ML</u>	<u>A078974 001</u>	Feb 18, 2011
<u>AA</u>	RUBICON	<u>250MG/5ML</u>	<u>A216492 001</u>	Jan 18, 2023
<u>AA</u>	TARO	<u>250MG/5ML</u>	<u>A076672 001</u>	Jul 03, 2013

NEURONTIN

<u>AA</u>	+! VIATRIS	<u>250MG/5ML</u>	<u>N021129 001</u>	Mar 02, 2000
-----------	------------	------------------	--------------------	--------------

TABLET;ORAL

GABAPENTIN

<u>AB1</u>	ACTAVIS ELIZABETH	<u>600MG</u>	<u>A075694 001</u>	Oct 21, 2004
<u>AB1</u>		<u>800MG</u>	<u>A075694 002</u>	Oct 21, 2004
<u>AB1</u>	ALKEM LABS LTD	<u>600MG</u>	<u>A206402 001</u>	Dec 23, 2015
<u>AB1</u>		<u>800MG</u>	<u>A206402 002</u>	Dec 23, 2015
<u>AB1</u>	APOTEX	<u>100MG</u>	<u>A077894 001</u>	Oct 10, 2006
<u>AB1</u>		<u>300MG</u>	<u>A077894 002</u>	Oct 10, 2006
<u>AB1</u>		<u>400MG</u>	<u>A077894 003</u>	Oct 10, 2006
<u>AB1</u>	ASCENT PHARMS INC	<u>600MG</u>	<u>A214957 001</u>	Oct 01, 2021
<u>AB1</u>		<u>800MG</u>	<u>A214957 002</u>	Oct 01, 2021
<u>AB1</u>	AUROBINDO PHARMA LTD	<u>600MG</u>	<u>A200651 001</u>	Oct 06, 2011
<u>AB1</u>		<u>800MG</u>	<u>A200651 002</u>	Oct 06, 2011
<u>AB1</u>	CSPC OUYI	<u>600MG</u>	<u>A207057 001</u>	Oct 26, 2017
<u>AB1</u>		<u>800MG</u>	<u>A207057 002</u>	Oct 26, 2017
<u>AB1</u>	GLENMARK PHARMS LTD	<u>600MG</u>	<u>A077662 001</u>	Aug 18, 2006
<u>AB1</u>		<u>800MG</u>	<u>A077662 002</u>	Aug 18, 2006
<u>AB1</u>	GRANULES	<u>600MG</u>	<u>A217116 001</u>	Mar 28, 2023
<u>AB1</u>		<u>800MG</u>	<u>A217116 002</u>	Mar 28, 2023
<u>AB1</u>	IVAX SUB TEVA PHARMS	<u>100MG</u>	<u>A076017 001</u>	Apr 28, 2004
<u>AB1</u>		<u>300MG</u>	<u>A076017 002</u>	Apr 28, 2004
<u>AB1</u>		<u>400MG</u>	<u>A076017 003</u>	Apr 28, 2004
<u>AB1</u>	LAURUS	<u>600MG</u>	<u>A217965 001</u>	Apr 08, 2024
<u>AB1</u>		<u>800MG</u>	<u>A217965 002</u>	Apr 08, 2024
<u>AB1</u>	RISING	<u>600MG</u>	<u>A217995 001</u>	Jul 19, 2023
<u>AB1</u>		<u>800MG</u>	<u>A217995 002</u>	Jul 19, 2023
<u>AB1</u>	RUBICON	<u>600MG</u>	<u>A077661 004</u>	Sep 13, 2006
<u>AB1</u>		<u>800MG</u>	<u>A077661 005</u>	Sep 13, 2006
<u>AB1</u>	SCIEGEN PHARMS INC	<u>600MG</u>	<u>A205101 001</u>	Feb 04, 2016
<u>AB1</u>		<u>800MG</u>	<u>A205101 002</u>	Feb 04, 2016
<u>AB1</u>	STRIDES PHARMA	<u>100MG</u>	<u>A211313 003</u>	Dec 20, 2024
<u>AB1</u>		<u>400MG</u>	<u>A211313 004</u>	Dec 20, 2024
<u>AB1</u>		<u>600MG</u>	<u>A203244 002</u>	Jul 12, 2013
<u>AB1</u>		<u>600MG</u>	<u>A211313 001</u>	Mar 04, 2024
<u>AB1</u>		<u>800MG</u>	<u>A203244 001</u>	Jul 12, 2013
<u>AB1</u>		<u>800MG</u>	<u>A211313 002</u>	Mar 04, 2024
<u>AB1</u>	ZYDUS PHARMS USA INC	<u>600MG</u>	<u>A078926 001</u>	Feb 11, 2011
<u>AB1</u>		<u>800MG</u>	<u>A078926 002</u>	Feb 11, 2011

NEURONTIN

<u>AB1</u>	+ VIATRIS	<u>600MG</u>	<u>N020882 001</u>	Oct 09, 1998
<u>AB1</u>	+!	<u>800MG</u>	<u>N020882 002</u>	Oct 09, 1998

GABAPENTIN

<u>AB2</u>	ABON PHARMS LLC	<u>600MG</u>	<u>A203643 001</u>	Sep 09, 2024
<u>AB2</u>	ANNORA PHARMA	<u>300MG</u>	<u>A218075 001</u>	Feb 26, 2024
<u>AB2</u>		<u>600MG</u>	<u>A218075 002</u>	Feb 26, 2024
<u>AB2</u>	HUMANWELL	<u>300MG</u>	<u>A216252 001</u>	Feb 26, 2024
<u>AB2</u>		<u>600MG</u>	<u>A216252 002</u>	Feb 26, 2024
<u>AB2</u>	ZYDUS PHARMS	<u>300MG</u>	<u>A203934 001</u>	Jan 24, 2024
<u>AB2</u>		<u>600MG</u>	<u>A203934 002</u>	Jan 24, 2024

GRALISE

<u>AB2</u>	+! ALMATICA	<u>300MG</u>	<u>N022544 001</u>	Jan 28, 2011
<u>AB2</u>	+	<u>600MG</u>	<u>N022544 002</u>	Jan 28, 2011
	+	450MG	N022544 003	Apr 18, 2023
	+	750MG	N022544 004	Apr 18, 2023
	+!	900MG	N022544 005	Apr 18, 2023

PRESCRIPTION DRUG PRODUCT LIST

GABAPENTIN ENACARBIL

TABLET, EXTENDED RELEASE;ORAL

HORIZANT

+	AZURITY	300MG	N022399	002	Dec 13, 2011
+	!	600MG	N022399	001	Apr 06, 2011

GADOBENATE DIMEGLUMINE

INJECTABLE; INTRAVENOUS

MULTIHANCE

+	BRACCO	2.645GM/5ML (529MG/ML)	N021357	001	Nov 23, 2004
+	!	5.29GM/10ML (529MG/ML)	N021357	002	Nov 23, 2004
+	!	7.935GM/15ML (529MG/ML)	N021357	003	Nov 23, 2004
+	!	10.58GM/20ML (529MG/ML)	N021357	004	Nov 23, 2004

MULTIHANCE MULTIPACK

+	BRACCO	26.45GM/50ML (529MG/ML)	N021358	001	Nov 23, 2004
+	!	52.9GM/100ML (529MG/ML)	N021358	002	Nov 23, 2004

GADOBUTROL

SOLUTION; INTRAVENOUS

GADAVIST

<u>AP</u>	+	BAYER HLTHCARE	<u>1.20944GM/2ML (604.72MG/ML)</u>	<u>N201277</u>	<u>006</u>	Dec 18, 2013
<u>AP</u>	+	!	<u>4.5354GM/7.5ML (604.72MG/ML)</u>	<u>N201277</u>	<u>001</u>	Mar 14, 2011
<u>AP</u>	+	!	<u>6.0472GM/10ML (604.72MG/ML)</u>	<u>N201277</u>	<u>002</u>	Mar 14, 2011
<u>AP</u>	+	!	<u>9.0708GM/15ML (604.72MG/ML)</u>	<u>N201277</u>	<u>003</u>	Mar 14, 2011
<u>AP</u>	+	!	<u>18.1416GM/30ML (604.72MG/ML)</u>	<u>N201277</u>	<u>004</u>	Mar 14, 2011
<u>AP</u>	+	!	<u>39.3068GM/65ML (604.72MG/ML)</u>	<u>N201277</u>	<u>005</u>	Mar 14, 2011

GADOBUTROL

<u>AP</u>		HAINAN POLY PHARM	<u>4.5354GM/7.5ML (604.72MG/ML)</u>	<u>A217480</u>	<u>001</u>	Mar 15, 2023
<u>AP</u>		!	<u>9.0708GM/15ML (604.72MG/ML)</u>	<u>A217480</u>	<u>002</u>	Mar 15, 2023
<u>AP</u>		HENGRUI PHARMA	<u>1.20944GM/2ML (604.72MG/ML)</u>	<u>A216081</u>	<u>003</u>	Nov 17, 2022
<u>AP</u>		!	<u>4.5354GM/7.5ML (604.72MG/ML)</u>	<u>A216081</u>	<u>004</u>	Nov 17, 2022
<u>AP</u>		!	<u>6.0472GM/10ML (604.72MG/ML)</u>	<u>A216081</u>	<u>005</u>	Nov 17, 2022
<u>AP</u>		!	<u>9.0708GM/15ML (604.72MG/ML)</u>	<u>A216081</u>	<u>006</u>	Nov 17, 2022
<u>AP</u>		!	<u>18.1416GM/30ML (604.72MG/ML)</u>	<u>A216081</u>	<u>001</u>	Sep 08, 2023
<u>AP</u>		!	<u>39.3068GM/65ML (604.72MG/ML)</u>	<u>A216081</u>	<u>002</u>	Sep 08, 2023

GADODIAMIDE

INJECTABLE; INJECTION

OMNISCAN

+	GE HEALTHCARE	287MG/ML	N020123	001	Jan 08, 1993
---	---------------	----------	---------	-----	--------------

GADOPICTENOL

SOLUTION; INTRAVENOUS

ELUCIREM

+	GUERBET	1.4553GM/3ML (485.1MG/ML)	N216986	001	Sep 21, 2022
+	!	3.63825GM/7.5ML (485.1MG/ML)	N216986	002	Sep 21, 2022
+	!	4.851GM/10ML (485.1MG/ML)	N216986	003	Sep 21, 2022
+	!	7.2765GM/15ML (485.1MG/ML)	N216986	004	Sep 21, 2022
+	!	14.553GM/30ML (485.1MG/ML)	N216986	005	Sep 21, 2022
+	!	24.255GM/50ML (485.1MG/ML)	N216986	006	Sep 21, 2022
+	!	48.51GM/100ML (485.1MG/ML)	N216986	007	Sep 21, 2022

GADOTERATE MEGLUMINE

SOLUTION; INTRAVENOUS

CLARISCAN

<u>AP</u>		GE HEALTHCARE	<u>7.538GM/20ML (376.9MG/ML)</u>	<u>A210016</u>	<u>003</u>	Nov 01, 2019
<u>AP</u>		!	<u>1.8845GM/5ML (376.9MG/ML)</u>	<u>A210016</u>	<u>005</u>	Nov 24, 2020
<u>AP</u>		!	<u>3.769GM/10ML (376.9MG/ML)</u>	<u>A210016</u>	<u>001</u>	Nov 01, 2019
<u>AP</u>		!	<u>5.6535GM/15ML (376.9MG/ML)</u>	<u>A210016</u>	<u>002</u>	Nov 01, 2019
<u>AP</u>		!	<u>37.69GM/100ML (376.9MG/ML)</u>	<u>A210016</u>	<u>004</u>	Aug 04, 2020

DOTAREM

<u>AP</u>	+	GUERBET	<u>37.69GM/100ML (376.9MG/ML)</u>	<u>N204781</u>	<u>001</u>	Mar 20, 2013
<u>AP</u>	+	!	<u>1.8845GM/5ML (376.9MG/ML)</u>	<u>N204781</u>	<u>005</u>	Mar 31, 2017
<u>AP</u>	+	!	<u>3.769GM/10ML (376.9MG/ML)</u>	<u>N204781</u>	<u>002</u>	Mar 20, 2013
<u>AP</u>	+	!	<u>5.6535GM/15ML (376.9MG/ML)</u>	<u>N204781</u>	<u>003</u>	Mar 20, 2013
<u>AP</u>	+	!	<u>7.538GM/20ML (376.9MG/ML)</u>	<u>N204781</u>	<u>004</u>	Mar 20, 2013

GADOTERATE MEGLUMINE

<u>AP</u>		HAINAN POLY	<u>1.8845GM/5ML (376.9MG/ML)</u>	<u>A218073</u>	<u>001</u>	Jun 17, 2024
<u>AP</u>		!	<u>3.769GM/10ML (376.9MG/ML)</u>	<u>A218073</u>	<u>002</u>	Jun 17, 2024
<u>AP</u>		!	<u>5.6535GM/15ML (376.9MG/ML)</u>	<u>A218073</u>	<u>003</u>	Jun 17, 2024
<u>AP</u>		!	<u>7.538GM/20ML (376.9MG/ML)</u>	<u>A218073</u>	<u>004</u>	Jun 17, 2024
<u>AP</u>		!	<u>37.69GM/100ML (376.9MG/ML)</u>	<u>A218073</u>	<u>005</u>	Jun 17, 2024
<u>AP</u>		HENGRUI PHARMA	<u>1.8845GM/5ML (376.9MG/ML)</u>	<u>A215304</u>	<u>001</u>	Apr 11, 2022
<u>AP</u>		!	<u>3.769GM/10ML (376.9MG/ML)</u>	<u>A215304</u>	<u>002</u>	Apr 11, 2022
<u>AP</u>		!	<u>5.6535GM/15ML (376.9MG/ML)</u>	<u>A215304</u>	<u>003</u>	Apr 11, 2022

PRESCRIPTION DRUG PRODUCT LIST

GADOTERATE MEGLUMINE

SOLUTION; INTRAVENOUS

GADOTERATE MEGLUMINE

<u>AP</u>		<u>7.538GM/20ML (376.9MG/ML)</u>	<u>A215304 004</u>	Apr 11, 2022
<u>AP</u>		<u>37.69GM/100ML (376.9MG/ML)</u>	<u>A215304 005</u>	Apr 15, 2022

GADOTERIDOL

INJECTABLE; INJECTION

PROHANCE

+	!	BRACCO	279.3MG/ML	N020131 001	Nov 16, 1992
---	---	--------	------------	-------------	--------------

PROHANCE MULTIPACK

+	!	BRACCO	279.3MG/ML	N021489 001	Oct 09, 2003
---	---	--------	------------	-------------	--------------

GADOXETATE DISODIUM

SOLUTION; INTRAVENOUS

EOVIST

+	!	BAYER HLTHCARE	1.8143GM/10ML (181.43MG/ML)	N022090 001	Jul 03, 2008
---	---	----------------	-----------------------------	-------------	--------------

+			2.72145GM/15ML (181.43MG/ML)	N022090 002	Feb 04, 2013
---	--	--	------------------------------	-------------	--------------

GALANTAMINE HYDROBROMIDE

CAPSULE, EXTENDED RELEASE; ORAL

GALANTAMINE HYDROBROMIDE

<u>AB</u>	!	AUROBINDO PHARMA	<u>EQ 8MG BASE</u>	<u>A204895 001</u>	Aug 05, 2016
<u>AB</u>			<u>EQ 16MG BASE</u>	<u>A204895 002</u>	Aug 05, 2016
<u>AB</u>			<u>EQ 24MG BASE</u>	<u>A204895 003</u>	Aug 05, 2016
<u>AB</u>		BARR	<u>EQ 8MG BASE</u>	<u>A078189 001</u>	Sep 15, 2008
<u>AB</u>			<u>EQ 16MG BASE</u>	<u>A078189 002</u>	Sep 15, 2008
<u>AB</u>			<u>EQ 24MG BASE</u>	<u>A078189 003</u>	Sep 15, 2008
<u>AB</u>		SUN PHARM	<u>EQ 8MG BASE</u>	<u>A090178 001</u>	Feb 02, 2011
<u>AB</u>			<u>EQ 16MG BASE</u>	<u>A090178 002</u>	Feb 02, 2011
<u>AB</u>			<u>EQ 24MG BASE</u>	<u>A090178 003</u>	Feb 02, 2011
<u>AB</u>		WATSON LABS	<u>EQ 8MG BASE</u>	<u>A079028 001</u>	Dec 15, 2008
<u>AB</u>			<u>EQ 16MG BASE</u>	<u>A079028 002</u>	Dec 15, 2008
<u>AB</u>			<u>EQ 24MG BASE</u>	<u>A079028 003</u>	Dec 15, 2008

SOLUTION; ORAL

GALANTAMINE HYDROBROMIDE

!		HIKMA	4MG/ML	A078185 001	Jan 30, 2009
---	--	-------	--------	-------------	--------------

TABLET; ORAL

GALANTAMINE HYDROBROMIDE

<u>AB</u>		AUROBINDO PHARMA LTD	<u>EQ 4MG BASE</u>	<u>A090957 001</u>	Mar 29, 2011
<u>AB</u>			<u>EQ 8MG BASE</u>	<u>A090957 002</u>	Mar 29, 2011
<u>AB</u>			<u>EQ 12MG BASE</u>	<u>A090957 003</u>	Mar 29, 2011
<u>AB</u>		BARR	<u>EQ 4MG BASE</u>	<u>A077605 001</u>	Aug 28, 2008
<u>AB</u>			<u>EQ 8MG BASE</u>	<u>A077605 002</u>	Aug 28, 2008
<u>AB</u>			<u>EQ 12MG BASE</u>	<u>A077605 003</u>	Aug 28, 2008
<u>AB</u>		DR REDDYS LABS LTD	<u>EQ 4MG BASE</u>	<u>A077593 001</u>	Sep 11, 2008
<u>AB</u>			<u>EQ 8MG BASE</u>	<u>A077593 002</u>	Sep 11, 2008
<u>AB</u>			<u>EQ 12MG BASE</u>	<u>A077593 003</u>	Sep 11, 2008
<u>AB</u>		SANDOZ	<u>EQ 4MG BASE</u>	<u>A077589 001</u>	Jun 22, 2009
<u>AB</u>			<u>EQ 8MG BASE</u>	<u>A077589 002</u>	Jun 22, 2009
<u>AB</u>			<u>EQ 12MG BASE</u>	<u>A077589 003</u>	Jun 22, 2009
<u>AB</u>	!	YABAO PHARM	<u>EQ 4MG BASE</u>	<u>A077604 001</u>	Feb 06, 2009
<u>AB</u>			<u>EQ 8MG BASE</u>	<u>A077604 002</u>	Feb 06, 2009
<u>AB</u>			<u>EQ 12MG BASE</u>	<u>A077604 003</u>	Feb 06, 2009
<u>AB</u>		ZYDUS PHARMS USA INC	<u>EQ 4MG BASE</u>	<u>A078898 001</u>	Feb 17, 2011
<u>AB</u>			<u>EQ 8MG BASE</u>	<u>A078898 002</u>	Feb 17, 2011
<u>AB</u>			<u>EQ 12MG BASE</u>	<u>A078898 003</u>	Feb 17, 2011

GALLIUM CITRATE GA-67

INJECTABLE; INJECTION

GALLIUM CITRATE GA 67

BS		CURIUM	2mCi/ML	N018058 001	
----	--	--------	---------	-------------	--

GALLIUM DOTATATE GA-68

POWDER; INTRAVENOUS

NETSPOT

+	!	AAA USA INC	2.1-5.5mCi/ML	N208547 001	Jun 01, 2016
---	---	-------------	---------------	-------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

GALLIUM GA-68 EDOTREOTIDE

SOLUTION; INTRAVENOUS

GALLIUM GA 68 EDOTREOTIDE

+! UIHC PET IMAGING 0.5-4mCi/ML N210828 001 Aug 21, 2019

GALLIUM GA-68 GOZETOTIDE

POWDER; INTRAVENOUS

ILLUCCIX

+! TELIX N/A N214032 001 Dec 17, 2021

LOCAMETZ

+! NOVARTIS N/A N215841 001 Mar 23, 2022

SOLUTION; INTRAVENOUS

GALLIUM GA 68 GOZETOTIDE

+! UNIV CA LOS ANGELES 0.5-5mCi/mL N212642 001 Dec 01, 2020

+! UNIV OF CA SAN FRAN 0.5-5mCi/mL N212643 001 Dec 01, 2020

GANAXOLONE

SUSPENSION; ORAL

ZTALMY

+! MARINUS 50MG/ML N215904 001 Jun 01, 2022

GANCICLOVIR

GEL; OPHTHALMIC

ZIRGAN

+! BAUSCH AND LOMB 0.15% N022211 001 Sep 15, 2009

SOLUTION; INTRAVENOUS

GANZYK-RTU

+! EXELA PHARMA 500MG/250ML (2MG/ML) N209347 001 Feb 17, 2017

GANCICLOVIR SODIUM

INJECTABLE; INJECTION

GANCICLOVIR SODIUM

AP	ENDO OPERATIONS	EQ 500MG BASE/VIAL	A204950 001	Dec 06, 2016
AP	FRESENIUS KABI USA	EQ 500MG BASE/VIAL	A090658 001	Jun 21, 2010
AP	HIKMA	EQ 500MG BASE/VIAL	A076222 001	Jul 16, 2003
AP	! PHARMASCIENCE INC	EQ 500MG BASE/VIAL	A207645 001	Dec 08, 2017
AP	SLATE RUN PHARMA	EQ 500MG BASE/VIAL	A204204 001	Nov 08, 2018

GANIRELIX ACETATE

INJECTABLE; INJECTION

FYREMADEL

AP	SUN PHARM	250MCG/0.5ML	A204246 001	Nov 30, 2018
-----------	-----------	---------------------	--------------------	--------------

GANIRELIX ACETATE

AP	AMPHASTAR PHARMS INC	250MCG/0.5ML	A212613 001	Apr 07, 2022
AP	GLAND PHARMA LTD	250MCG/0.5ML	A215658 001	Feb 28, 2023
AP	LUPIN LTD	250MCG/0.5ML	A216075 001	Nov 16, 2023
AP	MEITHEAL	250MCG/0.5ML	A214996 001	Jun 06, 2022
AP	+! ORGANON USA ORGANON	250MCG/0.5ML	N021057 001	Jul 29, 1999

GATIFLOXACIN

SOLUTION/DROPS; OPHTHALMIC

GATIFLOXACIN

AT	LUPIN LTD	0.5%	A202653 001	Aug 28, 2013
AT	SANDOZ	0.5%	A204227 001	Jul 11, 2016

ZYMAXID

AT	+! ABBVIE	0.5%	N022548 001	May 18, 2010
-----------	-----------	-------------	--------------------	--------------

GEFITINIB

TABLET; ORAL

GEFITINIB

AB	ACTAVIS LABS FL INC	250MG	A208913 001	Apr 26, 2023
AB	APOTEX	250MG	A209532 001	Sep 23, 2022
AB	NATCO	250MG	A212827 001	May 31, 2023
AB	QILU PHARM HAINAN	250MG	A211591 001	Feb 13, 2023

IRESSA

AB	+! ASTRAZENECA	250MG	N206995 001	Jul 13, 2015
-----------	----------------	--------------	--------------------	--------------

GEMCITABINE HYDROCHLORIDE

INJECTABLE; INJECTION

GEMCITABINE HYDROCHLORIDE

AP	DR REDDYS LABS LTD	EQ 200MG BASE/VIAL	A091365 001	Jul 25, 2011
AP		EQ 1GM BASE/VIAL	A091365 002	Jul 25, 2011
AP		EQ 2GM BASE/VIAL	A202997 001	May 07, 2013
AP	FRESENIUS KABI USA	EQ 200MG BASE/VIAL	A090799 001	Jul 25, 2011
AP		EQ 1GM BASE/VIAL	A090799 002	Jul 25, 2011
AP		EQ 2GM BASE/VIAL	A090242 003	May 16, 2011

PRESCRIPTION DRUG PRODUCT LIST

GEMCITABINE HYDROCHLORIDE

INJECTABLE; INJECTION

GEMCITABINE HYDROCHLORIDE

<u>AP</u>		<u>EQ 2GM BASE/VIAL</u>	<u>A090799 003</u>	May 16, 2011
<u>AP</u>	GLAND PHARMA LTD	<u>EQ 200MG BASE/VIAL</u>	<u>A204520 001</u>	Jan 05, 2016
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>A204520 002</u>	Jan 05, 2016
<u>AP</u>	HIKMA	<u>200MG/5.26ML (38MG/ML)</u>	<u>A213175 001</u>	Mar 07, 2023
<u>AP</u>		<u>1GM/26.3ML (38MG/ML)</u>	<u>A213175 002</u>	Mar 07, 2023
<u>AP</u>		<u>2GM/52.6ML (38MG/ML)</u>	<u>A213175 003</u>	Mar 07, 2023
<u>AP</u>	HIKMA INTL PHARMS	<u>EQ 200MG BASE/VIAL</u>	<u>A206617 001</u>	Jun 25, 2021
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>A206617 002</u>	Jun 25, 2021
<u>AP</u>	HOSPIRA	<u>EQ 200MG BASE/VIAL</u>	<u>A078339 001</u>	Jul 25, 2011
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>A078339 002</u>	Jul 25, 2011
<u>AP</u>	+! HOSPIRA INC	<u>200MG/5.26ML (38MG/ML)</u>	<u>N200795 001</u>	Aug 04, 2011
<u>AP</u>	+!	<u>1GM/26.3ML (38MG/ML)</u>	<u>N200795 002</u>	Aug 04, 2011
<u>AP</u>	!	<u>EQ 2GM BASE/VIAL</u>	<u>A079183 001</u>	Nov 15, 2010
<u>AP</u>	+!	<u>2GM/52.6ML (38MG/ML)</u>	<u>N200795 003</u>	Aug 04, 2011
<u>AP</u>	!	<u>EQ 200MG BASE/VIAL</u>	<u>A202485 001</u>	May 07, 2013
<u>AP</u>	!	<u>EQ 1GM BASE/VIAL</u>	<u>A202485 002</u>	May 07, 2013
<u>AP</u>	MEITHEAL	<u>200MG/5.26ML (38MG/ML)</u>	<u>A212129 001</u>	Dec 11, 2020
<u>AP</u>		<u>1GM/26.3ML (38MG/ML)</u>	<u>A212129 002</u>	Dec 11, 2020
<u>AP</u>		<u>2GM/52.6ML (38MG/ML)</u>	<u>A212129 003</u>	Dec 11, 2020
<u>AP</u>	MYLAN LABS LTD	<u>200MG/5.26ML (38MG/ML)</u>	<u>A205242 001</u>	Dec 06, 2017
<u>AP</u>		<u>1GM/26.3ML (38MG/ML)</u>	<u>A205242 002</u>	Dec 06, 2017
<u>AP</u>		<u>2GM/52.6ML (38MG/ML)</u>	<u>A205242 003</u>	Dec 06, 2017
<u>AP</u>	NOVAST LABS	<u>200MG/5.26ML (38MG/ML)</u>	<u>A210383 001</u>	Feb 14, 2019
<u>AP</u>		<u>1GM/26.3ML (38MG/ML)</u>	<u>A210383 002</u>	Feb 14, 2019
<u>AP</u>		<u>2GM/52.6ML (38MG/ML)</u>	<u>A210383 003</u>	Feb 14, 2019
<u>AP</u>	SAGENT PHARMS INC	<u>200MG/5.26ML (38MG/ML)</u>	<u>A209077 001</u>	Jul 20, 2018
<u>AP</u>		<u>1GM/26.3ML (38MG/ML)</u>	<u>A209077 002</u>	Jul 20, 2018
<u>AP</u>		<u>2GM/52.6ML (38MG/ML)</u>	<u>A209077 003</u>	Jul 20, 2018
<u>AP</u>	SHILPA	<u>EQ 200MG BASE/VIAL</u>	<u>A207575 001</u>	Feb 22, 2019
<u>AP</u>		<u>200MG/5.26ML (38MG/ML)</u>	<u>A210991 001</u>	Oct 04, 2019
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>A207575 002</u>	Feb 22, 2019
<u>AP</u>		<u>1GM/26.3ML (38MG/ML)</u>	<u>A210991 002</u>	Oct 04, 2019
<u>AP</u>		<u>2GM/52.6ML (38MG/ML)</u>	<u>A210991 003</u>	Oct 04, 2019
<u>AP</u>	SUN PHARM	<u>EQ 200MG BASE/VIAL</u>	<u>A078433 001</u>	Jul 25, 2011
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>A078433 002</u>	Jul 25, 2011
<u>AP</u>	TEYRO LABS	<u>EQ 200MG BASE/VIAL</u>	<u>A078759 001</u>	Jul 25, 2011
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>A078759 002</u>	Jul 25, 2011

SOLUTION; INTRAVENOUS

GEMCITABINE HYDROCHLORIDE

+!	ACCORD HLTHCARE	1GM/10ML (100MG/ML)	N209604 002	Aug 03, 2017
+!		1.5GM/15ML (100MG/ML)	N209604 003	Aug 03, 2017
+!		2GM/20ML (100MG/ML)	N209604 004	Aug 03, 2017
+!		200MG/2ML (100MG/ML)	N209604 001	Aug 03, 2017

GEMFIBROZIL

TABLET; ORAL

GEMFIBROZIL

<u>AB</u>	APOTEX	<u>600MG</u>	<u>A075034 001</u>	Jul 20, 1998
<u>AB</u>	ASCENT PHARMS INC	<u>600MG</u>	<u>A214603 001</u>	Jan 13, 2021
<u>AB</u>	AUROBINDO PHARMA LTD	<u>600MG</u>	<u>A202726 001</u>	Sep 16, 2015
<u>AB</u>	CADILA PHARMS LTD	<u>600MG</u>	<u>A203266 001</u>	Jun 17, 2016
<u>AB</u>	CHARTWELL MOLECULES	<u>600MG</u>	<u>A074270 001</u>	Sep 27, 1993
<u>AB</u>	EPIC PHARMA LLC	<u>600MG</u>	<u>A078012 001</u>	Mar 26, 2007
<u>AB</u>	IMPAX PHARMS	<u>600MG</u>	<u>A078207 001</u>	Jun 01, 2007
<u>AB</u>	INVAGEN PHARMS	<u>600MG</u>	<u>A077836 001</u>	Jul 27, 2006
<u>AB</u>	NORTHSTAR HLTHCARE	<u>600MG</u>	<u>A079072 001</u>	Sep 13, 2010
<u>AB</u>	+! PFIZER PHARMS	<u>600MG</u>	<u>N018422 003</u>	Nov 20, 1986

GENTAMICIN SULFATE

CREAM; TOPICAL

GENTAMICIN SULFATE

<u>AT</u>	COSETTE	<u>EQ 0.1% BASE</u>	<u>A064056 001</u>	Apr 29, 1994
<u>AT</u>	! PADAGIS US	<u>EQ 0.1% BASE</u>	<u>A062307 001</u>	

INJECTABLE; INJECTION

GENTAMICIN SULFATE

<u>AP</u>	EUGIA PHARMA	<u>EQ 10MG BASE/ML</u>	<u>A215236 001</u>	Jan 08, 2024
<u>AP</u>		<u>EQ 40MG BASE/ML</u>	<u>A215237 001</u>	Jan 08, 2024
<u>AP</u>	! FRESENIUS KABI USA	<u>EQ 10MG BASE/ML</u>	<u>A062366 002</u>	Feb 06, 1986

PRESCRIPTION DRUG PRODUCT LIST

GENTAMICIN SULFATE

INJECTABLE; INJECTION

GENTAMICIN SULFATE

<u>AP</u>	!		<u>EQ 40MG BASE/ML</u>	<u>A062366 001</u>	Aug 04, 1983
<u>AP</u>		HIKMA	<u>EQ 10MG BASE/ML</u>	<u>A062251 002</u>	
<u>AP</u>			<u>EQ 40MG BASE/ML</u>	<u>A062251 001</u>	
<u>AP</u>		HOSPIRA	<u>EQ 10MG BASE/ML</u>	<u>A062420 001</u>	Aug 15, 1983
<u>AP</u>			<u>EQ 40MG BASE/ML</u>	<u>A062420 002</u>	Aug 15, 1983

GENTAMICIN SULFATE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

<u>AP</u>		BAXTER HLTHCARE	<u>EQ 1.2MG BASE/ML</u>	<u>A062373 007</u>	Sep 07, 1982
<u>AP</u>			<u>EQ 1.6MG BASE/ML</u>	<u>A062373 008</u>	Sep 07, 1982
<u>AP</u>			<u>EQ 80MG BASE/100ML</u>	<u>A062373 002</u>	Sep 07, 1982
<u>AP</u>			<u>EQ 100MG BASE/100ML</u>	<u>A062373 005</u>	Sep 07, 1982
<u>AP</u>		HOSPIRA	<u>EQ 1.2MG BASE/ML</u>	<u>A062414 001</u>	Aug 15, 1983
<u>AP</u>			<u>EQ 1.6MG BASE/ML</u>	<u>A062414 003</u>	Aug 15, 1983
<u>AP</u>			<u>EQ 80MG BASE/100ML</u>	<u>A062414 008</u>	Aug 15, 1983
<u>AP</u>			<u>EQ 100MG BASE/100ML</u>	<u>A062414 010</u>	Aug 15, 1983
	!	BAXTER HLTHCARE	EQ 2MG BASE/ML	A062373 009	Sep 07, 1982
	!		EQ 120MG BASE/100ML	A062373 006	Sep 07, 1982

OINTMENT; OPHTHALMIC

GENTAMICIN SULFATE

! FERA PHARMS LLC

EQ 0.3% BASE

A065024 001 Jul 30, 2004

OINTMENT; TOPICAL

GENTAMICIN SULFATE

<u>AT</u>		COSETTE	<u>EQ 0.1% BASE</u>	<u>A064054 001</u>	Apr 29, 1994
<u>AT</u>		FOUGERA PHARMS INC	<u>EQ 0.1% BASE</u>	<u>A062533 001</u>	Oct 05, 1984
<u>AT</u>	!	PADAGIS US	<u>EQ 0.1% BASE</u>	<u>A062351 001</u>	Feb 18, 1982
<u>AT</u>		TARO	<u>EQ 0.1% BASE</u>	<u>A062477 001</u>	Dec 23, 1983

SOLUTION/DROPS; OPHTHALMIC

GENTAMICIN SULFATE

<u>AT</u>		BAUSCH AND LOMB	<u>EQ 0.3% BASE</u>	<u>A064048 001</u>	May 11, 1994
<u>AT</u>		PADAGIS US	<u>EQ 0.3% BASE</u>	<u>A065121 001</u>	Jan 30, 2004
<u>AT</u>	!	SANDOZ	<u>EQ 0.3% BASE</u>	<u>A062196 001</u>	

GILTERITINIB FUMARATE

TABLET; ORAL

XOSPATA

+! ASTELLAS

EQ 40MG BASE

N211349 001 Nov 28, 2018

GIVINOSTAT HYDROCHLORIDE

SUSPENSION; ORAL

DUVYZAT

+! ITALFARMACO SPA

EQ 8.86MG BASE/ML

N217865 001 Mar 21, 2024

GIVOSIRAN SODIUM

SOLUTION; SUBCUTANEOUS

GIVLAARI

+! ALNYLAM PHARMS INC

EQ 189MG BASE/ML (EQ 189MG BASE/ML)

N212194 001 Nov 20, 2019

GLASDEGIB MALEATE

TABLET; ORAL

DAURISMO

+ PFIZER

EQ 25MG BASE

N210656 001 Nov 21, 2018

+! EQ 100MG BASE

N210656 002 Nov 21, 2018

GLATIRAMER ACETATE

INJECTABLE; SUBCUTANEOUS

COPAXONE

<u>AP</u>	+!	TEVA PHARMS USA	<u>20MG/ML</u>	<u>N020622 002</u>	Feb 12, 2002
<u>AP</u>	+!		<u>40MG/ML</u>	<u>N020622 003</u>	Jan 28, 2014

GLATIRAMER ACETATE

<u>AP</u>		MYLAN	<u>20MG/ML</u>	<u>A091646 001</u>	Oct 03, 2017
<u>AP</u>			<u>40MG/ML</u>	<u>A206936 001</u>	Oct 03, 2017
<u>AP</u>		SYNTHON PHARMS INC	<u>20MG/ML</u>	<u>A203857 001</u>	Sep 25, 2024
<u>AP</u>			<u>40MG/ML</u>	<u>A206873 001</u>	Sep 25, 2024

GLATOPIA

<u>AP</u>		SANDOZ	<u>20MG/ML</u>	<u>A090218 001</u>	Apr 16, 2015
<u>AP</u>			<u>40MG/ML</u>	<u>A206921 001</u>	Feb 12, 2018

PRESCRIPTION DRUG PRODUCT LIST

GLECAPREVIR; PIBRENTASVIR

PELLETS; ORAL

MAVYRET

+! ABBVIE 50MG; 20MG/PACKET N215110 001 Jun 10, 2021

TABLET; ORAL

MAVYRET

+! ABBVIE 100MG; 40MG N209394 001 Aug 03, 2017

GLIMEPIRIDE

TABLET; ORAL

AMARYLAB +! SANOFI AVENTIS US 1MG N020496 001 Nov 30, 1995AB + 2MG N020496 002 Nov 30, 1995AB + 4MG N020496 003 Nov 30, 1995GLIMEPIRIDEAB ACCORD HLTHCARE 1MG A078181 001 Aug 23, 2007AB 2MG A078181 002 Aug 23, 2007AB 4MG A078181 003 Aug 23, 2007AB AUROBINDO PHARMA LTD 1MG A202759 001 Jun 29, 2012AB 2MG A202759 002 Jun 29, 2012AB 4MG A202759 003 Jun 29, 2012AB CARLSBAD 1MG A077911 001 Sep 22, 2009AB 2MG A077911 002 Sep 22, 2009AB 4MG A077911 003 Sep 22, 2009AB CHARTWELL MOLECULAR 1MG A077295 001 Oct 06, 2005AB 2MG A077295 002 Oct 06, 2005AB 4MG A077295 003 Oct 06, 2005AB DR REDDYS LABS LTD 1MG A077091 001 Oct 06, 2005AB 2MG A077091 002 Oct 06, 2005AB 4MG A077091 003 Oct 06, 2005AB INDOCO REMEDIES 1MG A202112 001 Apr 17, 2013AB 2MG A202112 002 Apr 17, 2013AB 4MG A202112 003 Apr 17, 2013AB MICRO LABS 1MG A091220 001 Jun 29, 2012AB 2MG A091220 002 Jun 29, 2012AB 4MG A091220 004 Jun 29, 2012AB 8MG A091220 006 Jun 29, 2012AB PRINSTON INC 1MG A077370 001 Dec 23, 2005AB 2MG A077370 002 Dec 23, 2005AB 4MG A077370 003 Dec 23, 2005AB 8MG A077370 004 Dec 23, 2005

MICRO LABS 3MG A091220 003 Jun 29, 2012

MICRO LABS 6MG A091220 005 Jun 29, 2012

GLIMEPIRIDE; PIOGLITAZONE HYDROCHLORIDE

TABLET; ORAL

DUETACTAB +! TAKEDA PHARMS USA 2MG; 30MG N021925 001 Jul 28, 2006AB + 4MG; 30MG N021925 002 Jul 28, 2006PIOGLITAZONE HYDROCHLORIDE AND GLIMEPIRIDEAB CHARTWELL RX 2MG; 30MG A201049 001 Jan 04, 2013AB 4MG; 30MG A201049 002 Jan 04, 2013GLIPIZIDE

TABLET; ORAL

GLIPIZIDEAB ANI PHARMS 5MG A074497 001 Aug 31, 1995AB 10MG A074497 002 Aug 31, 1995AB APOTEX 5MG A075795 001 Jun 13, 2001AB ! 10MG A075795 002 Jun 13, 2001AB AUROBINDO PHARMA USA 5MG A074226 001 May 10, 1994AB 10MG A074226 002 May 10, 1994AB RUBICON 5MG A214874 002 Oct 03, 2023AB 10MG A214874 003 Oct 03, 2023AB WATSON LABS TEVA 5MG A074223 001 Feb 27, 1995AB 10MG A074223 002 Feb 27, 1995

RUBICON 2.5MG A214874 001 Oct 03, 2023

TABLET, EXTENDED RELEASE; ORAL

GLIPIZIDEAB AUROBINDO PHARMA 2.5MG A206928 001 May 12, 2017AB 5MG A206928 002 May 12, 2017AB 10MG A206928 003 May 12, 2017

PRESCRIPTION DRUG PRODUCT LIST

GLIPIZIDE

TABLET, EXTENDED RELEASE;ORAL

GLIPIZIDE

<u>AB</u>	UNIQUE	<u>2.5MG</u>	<u>A204720 001</u>	Dec 29, 2016
<u>AB</u>		<u>5MG</u>	<u>A204720 002</u>	Dec 29, 2016
<u>AB</u>	!	<u>10MG</u>	<u>A204720 003</u>	Dec 29, 2016
<u>AB</u>	WATSON LABS	<u>2.5MG</u>	<u>A076467 003</u>	Mar 27, 2006
<u>AB</u>		<u>5MG</u>	<u>A076467 001</u>	Sep 08, 2003
<u>AB</u>		<u>10MG</u>	<u>A076467 002</u>	Nov 07, 2003
<u>AB</u>	ZYDUS PHARMS	<u>2.5MG</u>	<u>A203499 001</u>	Jul 16, 2018
<u>AB</u>		<u>5MG</u>	<u>A203499 002</u>	Jul 16, 2018
<u>AB</u>		<u>10MG</u>	<u>A203499 003</u>	Jul 16, 2018
<u>GLUCOTROL XL</u>				
<u>AB</u>	+ PFIZER	<u>2.5MG</u>	<u>N020329 003</u>	Aug 10, 1999
<u>AB</u>	+	<u>10MG</u>	<u>N020329 002</u>	Apr 26, 1994

GLIPIZIDE; METFORMIN HYDROCHLORIDE

TABLET;ORAL

GLIPIZIDE AND METFORMIN HYDROCHLORIDE

<u>AB</u>	EPIC PHARMA LLC	<u>2.5MG;250MG</u>	<u>A077507 001</u>	Oct 27, 2005
<u>AB</u>		<u>2.5MG;500MG</u>	<u>A077507 002</u>	Oct 27, 2005
<u>AB</u>		<u>5MG;500MG</u>	<u>A077507 003</u>	Oct 27, 2005
<u>AB</u>	HERITAGE	<u>2.5MG;250MG</u>	<u>A078728 001</u>	Jun 23, 2010
<u>AB</u>		<u>2.5MG;500MG</u>	<u>A078728 002</u>	Jun 23, 2010
<u>AB</u>		<u>5MG;500MG</u>	<u>A078728 003</u>	Jun 23, 2010
<u>AB</u>	TEVA PHARMS	<u>2.5MG;250MG</u>	<u>A077270 001</u>	Oct 28, 2005
<u>AB</u>		<u>2.5MG;500MG</u>	<u>A077270 002</u>	Oct 28, 2005
<u>AB</u>	!	<u>5MG;500MG</u>	<u>A077270 003</u>	Oct 28, 2005
<u>AB</u>	ZYDUS PHARMS USA INC	<u>2.5MG;250MG</u>	<u>A078905 001</u>	Jan 31, 2011
<u>AB</u>		<u>2.5MG;500MG</u>	<u>A078905 002</u>	Jan 31, 2011
<u>AB</u>		<u>5MG;500MG</u>	<u>A078905 003</u>	Jan 31, 2011

GLUCAGON

INJECTABLE; INJECTION

GLUCAGON

<u>AP</u>	! AMPHASTAR PHARMS INC	<u>1MG/VIAL</u>	<u>A208086 001</u>	Dec 28, 2020
<u>AP</u>	MYLAN LABS LTD	<u>1MG/VIAL</u>	<u>A204468 001</u>	Dec 12, 2024
POWDER;NASAL				
BAQSIMI				
	+! AMPHASTAR PHARMS INC	3MG	N210134 001	Jul 24, 2019
SOLUTION;SUBCUTANEOUS				
GVOKE HYOPEN				
	+! XERIS	0.5MG/0.1ML (0.5MG/0.1ML)	N212097 003	Sep 10, 2019
	+!	1MG/0.2ML (1MG/0.2ML)	N212097 004	Sep 10, 2019
GVOKE KIT				
	+! XERIS	1MG/0.2ML (1MG/0.2ML)	N212097 005	Aug 20, 2021
GVOKE PFS				
	+! XERIS	1MG/0.2ML (1MG/0.2ML)	N212097 002	Sep 10, 2019

GLUCAGON HYDROCHLORIDE

POWDER;INTRAMUSCULAR, INTRAVENOUS

GLUCAGON

+! FRESENIUS KABI USA EQ 1MG BASE/VIAL N201849 001 May 08, 2015

POWDER;INTRAMUSCULAR, INTRAVENOUS, SUBCUTANEOUS

GLUCAGON

+! FRESENIUS KABI USA EQ 1MG BASE/VIAL N201849 002 Sep 19, 2019

GLYBURIDE

TABLET;ORAL

GLYBURIDE (MICRONIZED)

<u>AB</u>	TEVA	<u>1.5MG</u>	<u>A074686 001</u>	Apr 20, 1999
<u>AB</u>		<u>3MG</u>	<u>A074686 002</u>	Apr 20, 1999
<u>AB</u>		<u>6MG</u>	<u>A074686 004</u>	Apr 20, 1999
<u>GLYNASE</u>				
<u>AB</u>	+ PFIZER	<u>1.5MG</u>	<u>N020051 001</u>	Mar 04, 1992
<u>AB</u>	+	<u>3MG</u>	<u>N020051 002</u>	Mar 04, 1992
<u>AB</u>	+	<u>6MG</u>	<u>N020051 004</u>	Sep 24, 1993

GLYBURIDE

<u>AB1</u>	CADILA PHARMS LTD	<u>1.25MG</u>	<u>A203379 001</u>	Jan 04, 2019
<u>AB1</u>		<u>2.5MG</u>	<u>A203379 002</u>	Jan 04, 2019
<u>AB1</u>		<u>5MG</u>	<u>A203379 003</u>	Jan 04, 2019
<u>AB1</u>	EPIC PHARMA LLC	<u>1.25MG</u>	<u>A076257 001</u>	Jun 27, 2002

PRESCRIPTION DRUG PRODUCT LIST

GLYBURIDE

TABLET; ORAL

GLYBURIDE

<u>AB1</u>		<u>2.5MG</u>	<u>A076257 002</u>	Jun 27, 2002
<u>AB1</u>		<u>5MG</u>	<u>A076257 003</u>	Jun 27, 2002
<u>AB1</u>	HERITAGE	<u>1.25MG</u>	<u>A090937 001</u>	Feb 28, 2011
<u>AB1</u>		<u>2.5MG</u>	<u>A090937 002</u>	Feb 28, 2011
<u>AB1</u>		<u>5MG</u>	<u>A090937 003</u>	Feb 28, 2011
<u>AB1</u>	ORIENT PHARMA CO LTD	<u>1.25MG</u>	<u>A206483 001</u>	Feb 22, 2019
<u>AB1</u>		<u>2.5MG</u>	<u>A206483 002</u>	Feb 22, 2019
<u>AB1</u>		<u>5MG</u>	<u>A206483 003</u>	Feb 22, 2019
<u>AB1</u>	TEVA	<u>1.25MG</u>	<u>A074388 001</u>	Aug 29, 1995
<u>AB1</u>		<u>2.5MG</u>	<u>A074388 002</u>	Aug 29, 1995
<u>AB1</u>	!	<u>5MG</u>	<u>A074388 003</u>	Aug 29, 1995
<u>AB1</u>	ZYDUS PHARMS	<u>1.25MG</u>	<u>A206749 001</u>	May 10, 2016
<u>AB1</u>		<u>2.5MG</u>	<u>A206749 002</u>	May 10, 2016
<u>AB1</u>		<u>5MG</u>	<u>A206749 003</u>	May 10, 2016

DIABETA

<u>AB2</u>	+	SANOFI AVENTIS US	<u>1.25MG</u>	<u>N017532 001</u>	May 01, 1984
<u>AB2</u>	+		<u>2.5MG</u>	<u>N017532 002</u>	May 01, 1984
<u>AB2</u>	+	!	<u>5MG</u>	<u>N017532 003</u>	May 01, 1984

GLYBURIDE

<u>AB2</u>		IMPAX LABS INC	<u>1.25MG</u>	<u>A206079 001</u>	Sep 30, 2015
<u>AB2</u>			<u>2.5MG</u>	<u>A206079 002</u>	Sep 30, 2015
<u>AB2</u>			<u>5MG</u>	<u>A206079 003</u>	Sep 30, 2015
		GLYBURIDE (MICRONIZED) TEVA	4.5MG	A074686 003	Apr 20, 1999

GLYBURIDE; METFORMIN HYDROCHLORIDE

TABLET; ORAL

GLYBURIDE AND METFORMIN HYDROCHLORIDE

<u>AB</u>		ACTAVIS ELIZABETH	<u>1.25MG; 250MG</u>	<u>A076716 001</u>	Jun 28, 2005
<u>AB</u>			<u>2.5MG; 500MG</u>	<u>A076716 002</u>	Jun 28, 2005
<u>AB</u>			<u>5MG; 500MG</u>	<u>A076716 003</u>	Jun 28, 2005
<u>AB</u>		AUROBINDO PHARMA	<u>1.25MG; 250MG</u>	<u>A077870 001</u>	Nov 14, 2007
<u>AB</u>	!		<u>2.5MG; 500MG</u>	<u>A077870 002</u>	Nov 14, 2007
<u>AB</u>			<u>5MG; 500MG</u>	<u>A077870 003</u>	Nov 14, 2007
<u>AB</u>		HERITAGE	<u>1.25MG; 250MG</u>	<u>A079009 001</u>	Jun 03, 2009
<u>AB</u>			<u>2.5MG; 500MG</u>	<u>A079009 002</u>	Jun 03, 2009
<u>AB</u>			<u>5MG; 500MG</u>	<u>A079009 003</u>	Jun 03, 2009
<u>AB</u>		IMPAX LABS INC	<u>1.25MG; 250MG</u>	<u>A076345 001</u>	Feb 18, 2004
<u>AB</u>			<u>2.5MG; 500MG</u>	<u>A076345 002</u>	Feb 18, 2004
<u>AB</u>			<u>5MG; 500MG</u>	<u>A076345 003</u>	Feb 18, 2004
<u>AB</u>		ZYDUS PHARMS	<u>1.25MG; 250MG</u>	<u>A206748 001</u>	Feb 29, 2016
<u>AB</u>			<u>2.5MG; 500MG</u>	<u>A206748 002</u>	Feb 29, 2016
<u>AB</u>			<u>5MG; 500MG</u>	<u>A206748 003</u>	Feb 29, 2016

GLYCEROL PHENYLBUTYRATE

LIQUID; ORAL

RAVICTI

	+	HORIZON THERAP US	1.1GM/ML	N203284 001	Feb 01, 2013
--	---	-------------------	----------	-------------	--------------

GLYCINE

SOLUTION; IRRIGATION

AMINOACETIC ACID 1.5% IN PLASTIC CONTAINER

<u>AT</u>	+	BAXTER HLTHCARE	<u>1.5GM/100ML</u>	<u>N017865 001</u>	
-----------	---	-----------------	--------------------	--------------------	--

GLYCINE 1.5% IN PLASTIC CONTAINER

<u>AT</u>		B BRAUN	<u>1.5GM/100ML</u>	<u>N016784 001</u>	
<u>AT</u>		ICU MEDICAL INC	<u>1.5GM/100ML</u>	<u>N018315 001</u>	

GLYCOPYRROLATE

INJECTABLE; INJECTION

GLYCOPYRROLATE

<u>AP</u>		ACCORD HLTHCARE	<u>0.2MG/ML</u>	<u>A213238 001</u>	Jul 08, 2020
<u>AP</u>		ALEMBIC	<u>0.2MG/ML</u>	<u>A214635 001</u>	Oct 26, 2022
<u>AP</u>		AM REGENT	<u>0.2MG/ML</u>	<u>A089335 001</u>	Jul 23, 1986
<u>AP</u>		AMNEAL	<u>0.2MG/ML</u>	<u>A208973 001</u>	Jun 15, 2017
<u>AP</u>			<u>0.2MG/ML</u>	<u>A215333 001</u>	Oct 21, 2022
<u>AP</u>		APOTEX	<u>0.2MG/ML</u>	<u>A210246 001</u>	Oct 29, 2019
<u>AP</u>		CAPLIN	<u>0.2MG/ML</u>	<u>A211705 001</u>	Mar 20, 2019
<u>AP</u>		FRESENIUS KABI USA	<u>0.2MG/ML</u>	<u>A209024 001</u>	Oct 31, 2018
<u>AP</u>			<u>0.2MG/ML</u>	<u>A209328 001</u>	Oct 27, 2017
<u>AP</u>		GLAND PHARMA LTD	<u>0.2MG/ML</u>	<u>A212612 001</u>	Sep 30, 2019
<u>AP</u>	!	HIKMA FARMACEUTICA	<u>0.2MG/ML</u>	<u>A090963 001</u>	Sep 21, 2011

PRESCRIPTION DRUG PRODUCT LIST

GLYCOPYRROLATE

INJECTABLE; INJECTION

GLYCOPYRROLATE

<u>AP</u>	MEITHEAL	<u>0.2MG/ML</u>	<u>A212802</u>	<u>001</u>	Jul 06, 2021
<u>AP</u>	OMNIVIUM PHARMS	<u>0.2MG/ML</u>	<u>A212871</u>	<u>001</u>	Nov 29, 2022
<u>AP</u>	PIRAMAL CRITICAL	<u>0.2MG/ML</u>	<u>A210842</u>	<u>001</u>	Oct 25, 2018
<u>AP</u>	PRINSTON INC	<u>0.2MG/ML</u>	<u>A210927</u>	<u>001</u>	Oct 31, 2018
<u>AP</u>	SAGENT	<u>0.2MG/ML</u>	<u>A210083</u>	<u>001</u>	Feb 21, 2020
<u>AP</u>	SANDOZ	<u>0.2MG/ML</u>	<u>A211334</u>	<u>001</u>	May 14, 2019
<u>AP</u>	SOMERSET THERAPS LLC	<u>0.2MG/ML</u>	<u>A207639</u>	<u>001</u>	Jun 23, 2017
<u>AP</u>	UMEDICA	<u>0.2MG/ML</u>	<u>A212591</u>	<u>001</u>	Oct 13, 2021
<u>AP</u>	XIROMED	<u>0.2MG/ML</u>	<u>A212227</u>	<u>001</u>	Mar 04, 2021

SOLUTION; INTRAMUSCULAR, INTRAVENOUS

GLYCOPYRROLATE

+! FRESENIUS KABI USA 0.6MG/3ML (0.2MG/ML) N214919 001 Apr 21, 2022

GLYRX-PF

+! EXELA PHARMA 0.2MG/ML (0.2MG/ML) N210997 001 Jul 11, 2018

+! 0.4MG/2ML (0.2MG/ML) N210997 002 Jul 11, 2018

+! 1MG/5ML (0.2MG/ML) N210997 003 Apr 09, 2020

SOLUTION; ORAL

CUVPOSAAA +! MERZ PHARMS 1MG/5ML N022571 001 Jul 28, 2010GLYCOPYRROLATEAA ANNORA PHARMA 1MG/5ML A213698 001 Jul 05, 2022AA AUROBINDO PHARMA
LTD 1MG/5ML A214847 001 Dec 31, 2024AA CHARTWELL RX 1MG/5ML A216368 001 Aug 28, 2024AA ENDO OPERATIONS 1MG/5ML A204438 001 Aug 09, 2021AA GRANULES 1MG/5ML A214735 001 Aug 19, 2024AA SAPTALIS PHARMS 1MG/5ML A216297 001 Oct 07, 2024AA SUVEN PHARMS 1MG/5ML A212467 001 Jul 05, 2022

TABLET; ORAL

GLYCOPYRROLATEAA ADAPTIS 1MG A091182 001 Feb 03, 2014AA 2MG A091182 002 Feb 03, 2014AA ALEMBIC 1MG A203657 001 Nov 30, 2018AA 2MG A203657 002 Nov 30, 2018AA APPCO 1MG A207201 001 Jan 03, 2017AA 2MG A207201 002 Jan 03, 2017AA AUROBINDO PHARMA 1MG A202675 001 Apr 15, 2013AA 2MG A202675 002 Oct 30, 2018AA ! ENDO OPERATIONS 1MG A040653 001 Aug 31, 2006AA ! 2MG A040653 002 Aug 31, 2006AA INDICUS PHARMA 1MG A040847 001 Mar 21, 2008AA 2MG A040847 002 Mar 21, 2008AA NATCO 1MG A091413 001 Jun 20, 2016AA 2MG A091413 002 Jun 20, 2016AA OXFORD PHARMS 1MG A090020 001 Oct 19, 2011AA 2MG A090020 002 Oct 19, 2011AA QUAGEN 1MG A212696 001 Jul 10, 2024AA 2MG A212696 002 Jul 10, 2024AA RISING 1MG A040821 001 Dec 29, 2008AA 2MG A040821 002 Dec 29, 2008AA SUN PHARM INDS LTD 1MG A040844 001 Aug 18, 2009AA 2MG A040844 002 Aug 18, 2009AA VELZEN PHARMA PVT 1MG A090195 001 Sep 21, 2012AA 2MG A090195 002 Sep 21, 2012ROBINULAA + CASPER PHARMA LLC 1MG N012827 001ROBINUL FORTEAA + CASPER PHARMA LLC 2MG N012827 002GLYCOPYRROLATE

! LGM PHARMA 1.5MG A091522 001 Mar 12, 2012

GLYCOPYRROLATE; NEOSTIGMINE METHYLSULFATE

SOLUTION; INTRAVENOUS

PREVDUO

+! AZURITY 0.6MG/3ML (0.2MG/ML); 3MG/3ML (1MG/ML) N216903 001 Feb 23, 2023

PRESCRIPTION DRUG PRODUCT LIST

GLYCOPYRRONIUM TOSYLATE

CLOTH; TOPICAL

QBREXZA

+! JOURNEY

EQ 2.4% BASE

N210361 001 Jun 28, 2018

GOLODIRSEN

SOLUTION; INTRAVENOUS

VYONDYS 53

+! SAREPTA THERAPS INC 100MG/2ML (50MG/ML)

N211970 001 Dec 12, 2019

GOSERELIN ACETATE

IMPLANT; IMPLANTATION

ZOLADEX

+! TERSERA

EQ 3.6MG BASE

N019726 001 Dec 29, 1989

+!

EQ 10.8MG BASE

N020578 001 Jan 11, 1996

GRAMICIDIN; NEOMYCIN SULFATE; POLYMYXIN B SULFATE

SOLUTION/DROPS; OPHTHALMIC

NEOMYCIN AND POLYMYXIN B SULFATES AND GRAMICIDIN**AT** ! BAUSCH AND LOMB 0.025MG/ML;EQ 1.75MG BASE/ML;10,000 **A064047 001** Jan 31, 1996**AT** NORDIC PHARMA 0.025MG/ML;EQ 1.75MG BASE/ML;10,000 **A065187 001** Oct 28, 2005
UNITS/MLNEOSPORIN**AT** ! MONARCH PHARMS 0.025MG/ML;EQ 1.75MG BASE/ML;10,000 **A060582 001**
UNITS/MLGRANISETRON

FILM, EXTENDED RELEASE; TRANSDERMAL

SANCUSO

+! CUMBERLAND

3.1MG/24HR

N022198 001 Sep 12, 2008

INJECTABLE; SUBCUTANEOUS

SUSTOL

+! HERON THERAPS INC 10MG/0.4ML (10MG/0.4ML)

N022445 001 Aug 09, 2016

GRANISETRON HYDROCHLORIDE

INJECTABLE; INJECTION

GRANISETRON HYDROCHLORIDE**AP** AMNEAL EQ 0.1MG BASE/ML (EQ 0.1MG BASE/ML) **A078262 001** Dec 31, 2007**AP** EQ 1MG BASE/ML (EQ 1MG BASE/ML) **A078258 001** Jun 30, 2008**AP** EQ 4MG BASE/4ML (EQ 1MG BASE/ML) **A078258 002** Jun 30, 2008**AP** BIONPHARMA EQ 0.1MG BASE/ML (EQ 0.1MG BASE/ML) **A078863 001** Jun 30, 2008**AP** EQ 4MG BASE/4ML (EQ 1MG BASE/ML) **A078880 001** Jun 30, 2008**AP** ! DR REDDYS EQ 0.1MG BASE/ML (EQ 0.1MG BASE/ML) **A078392 001** Dec 31, 2007**AP** ! EQ 4MG BASE/4ML (EQ 1MG BASE/ML) **A077297 001** Jun 30, 2008**AP** FRESENIUS KABI USA EQ 0.1MG BASE/ML (EQ 0.1MG BASE/ML) **A078522 001** Dec 31, 2007**AP** EQ 4MG BASE/4ML (EQ 1MG BASE/ML) **A078090 001** Jun 30, 2008**AP** HIKMA EQ 0.1MG BASE/ML (EQ 0.1MG BASE/ML) **A077913 001** Jun 26, 2008**AP** EQ 4MG BASE/4ML (EQ 1MG BASE/ML) **A077186 001** Jun 30, 2008**AP** EQ 1MG BASE/ML (EQ 1MG BASE/ML) **A077187 001** Jun 30, 2008**AP** EQ 4MG BASE/4ML (EQ 1MG BASE/ML) **A077177 001** Dec 31, 2007**AP** HIKMA FARMACEUTICA EQ 1MG BASE/ML (EQ 1MG BASE/ML) **A078629 001** Dec 23, 2009**AP** EQ 4MG BASE/4ML (EQ 1MG BASE/ML) **A078629 002** Dec 23, 2009**AP** MYLAN ASI EQ 0.1MG BASE/ML (EQ 0.1MG BASE/ML) **A091136 001** Apr 09, 2010**AP** EQ 1MG BASE/ML (EQ 1MG BASE/ML) **A091136 002** Apr 09, 2010**AP** EQ 4MG BASE/4ML (EQ 1MG BASE/ML) **A091137 002** Apr 09, 2010**AP** RISING EQ 1MG BASE/ML (EQ 1MG BASE/ML) **A079078 001** Sep 14, 2009**AP** EQ 4MG BASE/4ML (EQ 1MG BASE/ML) **A079078 002** Sep 14, 2009**AP** SANDOZ EQ 0.1MG BASE/ML (EQ 0.1MG BASE/ML) **A078534 001** Apr 30, 2009**AP** EQ 1MG BASE/ML (EQ 1MG BASE/ML) **A078531 001** Apr 30, 2009**AP** EQ 4MG BASE/4ML (EQ 1MG BASE/ML) **A078531 002** Apr 30, 2009**AP** SANDOZ INC EQ 1MG BASE/ML (EQ 1MG BASE/ML) **A078835 001** Jun 30, 2008**AP** EQ 4MG BASE/4ML (EQ 1MG BASE/ML) **A078835 002** Jun 30, 2008GRANISETRON HYDROCHLORIDE PRESERVATIVE FREE**AP** BIONPHARMA EQ 1MG BASE/ML (EQ 1MG BASE/ML) **A078863 002** Jun 30, 2008**AP** ! FRESENIUS KABI USA EQ 1MG BASE/ML (EQ 1MG BASE/ML) **A078096 001** Jun 30, 2008

TABLET; ORAL

GRANISETRON HYDROCHLORIDE**AB** APOTEX INC EQ 1MG BASE **A078843 001** Feb 27, 2008**AB** CHARTWELL MOLECULAR EQ 1MG BASE **A078037 001** Feb 27, 2008**AB** DR REDDYS LABS LTD EQ 1MG BASE **A078846 001** Feb 27, 2009**AB** NATCO PHARMA EQ 1MG BASE **A078969 001** Jun 22, 2009**AB** ! ORBION PHARMS EQ 1MG BASE **A078678 001** Feb 13, 2008**AB** TARO EQ 1MG BASE **A090817 001** May 28, 2010

PRESCRIPTION DRUG PRODUCT LIST

GRISEOFULVIN, MICROSIZE

SUSPENSION; ORAL

GRISEOFULVIN

AB	!	ACTAVIS MID ATLANTIC	125MG/5ML	A065394 001	Jul 06, 2007
AB		CHARTWELL RX	125MG/5ML	A065200 001	Mar 02, 2005
AB		CIPLA	125MG/5ML	A065354 001	Sep 10, 2007
AB		COSETTE	125MG/5ML	A065438 001	Oct 08, 2010

TABLET; ORAL

FULVICIN-U/F

AB		CHARTWELL RX	250MG	A060569 002	
AB			500MG	A060569 001	
<u>GRISEOFULVIN</u>					
AB		SANDOZ	250MG	A091592 001	Aug 07, 2013
AB	!		500MG	A091592 002	Aug 07, 2013
AB		SIGMAPHARM LABS LLC	500MG	A202482 001	Oct 22, 2012

GRISEOFULVIN, ULTRAMICROSIZE

TABLET; ORAL

FULVICIN P/G

AB		CHARTWELL RX	125MG	A061996 001	
AB			250MG	A061996 002	
<u>GRIS-PEG</u>					
AB	+	BAUSCH	125MG	N050475 001	
AB	+		250MG	N050475 002	

GRISEOFULVIN, ULTRAMICROSIZE

AB		MOUNTAIN	125MG	A204371 001	Jan 09, 2014
AB	!		250MG	A204371 002	Jan 09, 2014
AB		SANDOZ	125MG	A202805 001	Dec 26, 2018
AB			250MG	A202805 002	Dec 26, 2018

GRISEOFULVIN, ULTRAMICROSIZE

AB		SIGMAPHARM LABS LLC	125MG	A202545 001	Oct 22, 2012
AB			250MG	A202545 002	Oct 22, 2012
		FULVICIN P/G 165			
		CHARTWELL RX	165MG	A061996 003	Apr 06, 1982
		FULVICIN P/G 330			
		CHARTWELL RX	330MG	A061996 004	Apr 06, 1982

GUANFACINE HYDROCHLORIDE

TABLET; ORAL

GUANFACINE HYDROCHLORIDE

AB		AJANTA PHARMA LTD	EQ 1MG BASE	A217293 001	Apr 27, 2023
AB			EQ 2MG BASE	A217293 002	Apr 27, 2023
AB		AMNEAL PHARM	EQ 1MG BASE	A075109 001	Nov 25, 1998
AB	!		EQ 2MG BASE	A075109 002	Nov 25, 1998
AB		AUROBINDO PHARMA USA	EQ 1MG BASE	A074796 001	Jan 27, 1997
AB			EQ 2MG BASE	A074796 002	Jan 27, 1997
AB		EPIC PHARMA LLC	EQ 1MG BASE	A074673 001	Feb 28, 1997
AB			EQ 2MG BASE	A074673 002	Feb 28, 1997
AB		I 3 PHARMS	EQ 1MG BASE	A216828 001	Oct 05, 2023
AB			EQ 2MG BASE	A216828 002	Oct 05, 2023
AB		RUBICON	EQ 1MG BASE	A216762 001	Oct 17, 2023
AB			EQ 2MG BASE	A216762 002	Oct 17, 2023
AB		TWI PHARMS	EQ 1MG BASE	A216399 001	Jun 08, 2022
AB			EQ 2MG BASE	A216399 002	Jun 08, 2022
AB		UNICHEM	EQ 1MG BASE	A214689 001	Mar 03, 2021
AB			EQ 2MG BASE	A214689 002	Mar 03, 2021
AB		WATSON LABS	EQ 1MG BASE	A074145 001	Oct 17, 1995
AB			EQ 2MG BASE	A074145 002	Oct 17, 1995
AB		XIROMED	EQ 1MG BASE	A218326 001	Feb 23, 2024
AB			EQ 2MG BASE	A218326 002	Feb 23, 2024

TABLET, EXTENDED RELEASE; ORAL

GUANFACINE HYDROCHLORIDE

AB		ACTAVIS ELIZABETH	EQ 1MG BASE	A200881 001	Oct 05, 2012
AB			EQ 2MG BASE	A200881 002	Oct 05, 2012
AB			EQ 3MG BASE	A200881 003	Oct 05, 2012
AB			EQ 4MG BASE	A200881 004	Oct 05, 2012
AB		ALEMBIC	EQ 1MG BASE	A217269 001	Aug 07, 2023
AB			EQ 2MG BASE	A217269 002	Aug 07, 2023
AB			EQ 3MG BASE	A217269 003	Aug 07, 2023
AB			EQ 4MG BASE	A217269 004	Aug 07, 2023
AB		APOTEX	EQ 1MG BASE	A205430 001	Jul 25, 2018
AB			EQ 2MG BASE	A205430 002	Jul 25, 2018

PRESCRIPTION DRUG PRODUCT LIST

GUANFACINE HYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL

GUANFACINE HYDROCHLORIDE

<u>AB</u>		<u>EQ 3MG BASE</u>	<u>A205430 003</u>	Jul 25, 2018
<u>AB</u>		<u>EQ 4MG BASE</u>	<u>A205430 004</u>	Jul 25, 2018
<u>AB</u>	SANDOZ	<u>EQ 1MG BASE</u>	<u>A202568 001</u>	Jun 03, 2015
<u>AB</u>		<u>EQ 2MG BASE</u>	<u>A202568 002</u>	Jun 03, 2015
<u>AB</u>		<u>EQ 3MG BASE</u>	<u>A202568 003</u>	Jun 03, 2015
<u>AB</u>		<u>EQ 4MG BASE</u>	<u>A202568 004</u>	Jun 03, 2015
<u>AB</u>	SUN PHARM	<u>EQ 1MG BASE</u>	<u>A205689 001</u>	Nov 16, 2017
<u>AB</u>		<u>EQ 2MG BASE</u>	<u>A205689 002</u>	Nov 16, 2017
<u>AB</u>		<u>EQ 3MG BASE</u>	<u>A205689 003</u>	Nov 16, 2017
<u>AB</u>		<u>EQ 4MG BASE</u>	<u>A205689 004</u>	Nov 16, 2017
<u>AB</u>	TEVA PHARMS USA	<u>EQ 1MG BASE</u>	<u>A201382 001</u>	Jun 02, 2015
<u>AB</u>		<u>EQ 2MG BASE</u>	<u>A201382 002</u>	Jun 02, 2015
<u>AB</u>		<u>EQ 3MG BASE</u>	<u>A201382 003</u>	Jun 02, 2015
<u>AB</u>		<u>EQ 4MG BASE</u>	<u>A201382 004</u>	Jun 02, 2015
<u>AB</u>	TWI PHARMS	<u>EQ 1MG BASE</u>	<u>A201408 001</u>	Jun 02, 2015
<u>AB</u>		<u>EQ 2MG BASE</u>	<u>A201408 002</u>	Jun 02, 2015
<u>AB</u>		<u>EQ 3MG BASE</u>	<u>A201408 003</u>	Jun 02, 2015
<u>AB</u>		<u>EQ 4MG BASE</u>	<u>A201408 004</u>	Jun 02, 2015
<u>AB</u>	UNICHEM	<u>EQ 1MG BASE</u>	<u>A219033 001</u>	Nov 19, 2024
<u>AB</u>		<u>EQ 2MG BASE</u>	<u>A219033 002</u>	Nov 19, 2024
<u>AB</u>		<u>EQ 3MG BASE</u>	<u>A219033 003</u>	Nov 19, 2024
<u>AB</u>		<u>EQ 4MG BASE</u>	<u>A219033 004</u>	Nov 19, 2024
<u>AB</u>	YICHANG HUMANWELL	<u>EQ 1MG BASE</u>	<u>A213428 001</u>	Nov 25, 2020
<u>AB</u>		<u>EQ 2MG BASE</u>	<u>A213428 002</u>	Nov 25, 2020
<u>AB</u>		<u>EQ 3MG BASE</u>	<u>A213428 003</u>	Nov 25, 2020
<u>AB</u>		<u>EQ 4MG BASE</u>	<u>A213428 004</u>	Nov 25, 2020
<u>INTUNIV</u>				
<u>AB</u>	+	TAKEDA PHARMS USA	<u>EQ 1MG BASE</u>	<u>N022037 001</u> Sep 02, 2009
<u>AB</u>	+		<u>EQ 2MG BASE</u>	<u>N022037 002</u> Sep 02, 2009
<u>AB</u>	+		<u>EQ 3MG BASE</u>	<u>N022037 003</u> Sep 02, 2009
<u>AB</u>	+		<u>EQ 4MG BASE</u>	<u>N022037 004</u> Sep 02, 2009

HALCINONIDE

CREAM;TOPICAL

HALCINONIDE

<u>AB</u>	CHARTWELL RX	<u>0.1%</u>	<u>A214723 001</u>	Sep 08, 2021
<u>AB</u>	MYLAN	<u>0.1%</u>	<u>A211027 001</u>	Aug 12, 2019
<u>HALOG</u>				
<u>AB</u>	+	SUN PHARM INDS INC	<u>0.1%</u>	<u>N017556 001</u>
SOLUTION;TOPICAL				
HALCINONIDE				
	ENCUBE	0.1%	A217671 001	May 29, 2024

HALOBETASOL PROPIONATE

AEROSOL, FOAM;TOPICAL

HALOBETASOL PROPIONATE

<u>AB</u>	PADAGIS ISRAEL	<u>0.05%</u>	<u>A215266 001</u>	Aug 11, 2023
<u>LEXETTE</u>				
<u>AB</u>	+	MAYNE PHARMA	<u>0.05%</u>	<u>N210566 001</u> May 24, 2018
CREAM;TOPICAL				
<u>HALOBETASOL PROPIONATE</u>				
<u>AB</u>	!	COSETTE	<u>0.05%</u>	<u>A078162 001</u> Apr 24, 2007
<u>AB</u>		FOUGERA PHARMS	<u>0.05%</u>	<u>A077001 001</u> Dec 16, 2004
<u>AB</u>		PADAGIS ISRAEL	<u>0.05%</u>	<u>A077123 001</u> Dec 16, 2004
<u>AB</u>		TARO	<u>0.05%</u>	<u>A077227 001</u> Aug 04, 2005
LOTION;TOPICAL				
BRYHALI				
	+	BAUSCH	0.01%	N209355 001 Nov 06, 2018
OINTMENT;TOPICAL				
<u>HALOBETASOL PROPIONATE</u>				
<u>AB</u>		COSETTE	<u>0.05%</u>	<u>A077109 001</u> Jun 14, 2005
<u>AB</u>	!	PADAGIS ISRAEL	<u>0.05%</u>	<u>A076872 001</u> Dec 16, 2004
<u>AB</u>		QUAGEN	<u>0.05%</u>	<u>A213560 001</u> Oct 06, 2020
<u>AB</u>		TARO	<u>0.05%</u>	<u>A076994 001</u> Dec 16, 2004

PRESCRIPTION DRUG PRODUCT LIST

HALOBETASOL PROPIONATE; TAZAROTENE

LOTION; TOPICAL

DUOBRII

+! BAUSCH 0.01%; 0.045% N209354 001 Apr 25, 2019

HALOPERIDOL

TABLET; ORAL

HALOPERIDOL

<u>AB</u>	AIPING PHARM INC	<u>2MG</u>	<u>A071130 001</u>	Feb 17, 1987
<u>AB</u>		<u>5MG</u>	<u>A071130 002</u>	Feb 17, 1987
<u>AB</u>		<u>10MG</u>	<u>A071130 003</u>	May 12, 1987
<u>AB</u>	AUROBINDO PHARMA LTD	<u>0.5MG</u>	<u>A218789 001</u>	Apr 19, 2024
<u>AB</u>		<u>1MG</u>	<u>A218789 002</u>	Apr 19, 2024
<u>AB</u>		<u>2MG</u>	<u>A218789 003</u>	Apr 19, 2024
<u>AB</u>		<u>5MG</u>	<u>A218789 004</u>	Apr 19, 2024
<u>AB</u>		<u>10MG</u>	<u>A218789 005</u>	Apr 19, 2024
<u>AB</u>		<u>20MG</u>	<u>A218789 006</u>	Apr 19, 2024
<u>AB</u>	CHARTWELL RX	<u>0.5MG</u>	<u>A071209 002</u>	Nov 17, 1986
<u>AB</u>		<u>1MG</u>	<u>A071209 003</u>	Nov 17, 1986
<u>AB</u>		<u>5MG</u>	<u>A071209 001</u>	Nov 17, 1986
<u>AB</u>		<u>10MG</u>	<u>A071210 001</u>	Mar 11, 1988
<u>AB</u>		<u>20MG</u>	<u>A071211 001</u>	Mar 11, 1988
<u>AB</u>	INNOGENIX	<u>0.5MG</u>	<u>A071173 002</u>	Jan 02, 1987
<u>AB</u>		<u>1MG</u>	<u>A071173 003</u>	Jan 02, 1987
<u>AB</u>		<u>2MG</u>	<u>A071173 004</u>	Jan 02, 1987
<u>AB</u>		<u>5MG</u>	<u>A071173 005</u>	Jan 07, 1988
<u>AB</u>		<u>10MG</u>	<u>A071173 001</u>	Jan 07, 1988
<u>AB</u>		<u>20MG</u>	<u>A071173 006</u>	Jan 07, 1988
<u>AB</u>	MANKIND PHARMA	<u>0.5MG</u>	<u>A216918 001</u>	Apr 11, 2024
<u>AB</u>		<u>1MG</u>	<u>A216918 002</u>	Apr 11, 2024
<u>AB</u>		<u>2MG</u>	<u>A216918 003</u>	Apr 11, 2024
<u>AB</u>		<u>5MG</u>	<u>A216918 004</u>	Apr 11, 2024
<u>AB</u>		<u>10MG</u>	<u>A216918 005</u>	Apr 11, 2024
<u>AB</u>		<u>20MG</u>	<u>A216918 006</u>	Apr 11, 2024
<u>AB</u>	MSN	<u>0.5MG</u>	<u>A216004 001</u>	Nov 18, 2022
<u>AB</u>		<u>1MG</u>	<u>A216004 002</u>	Nov 18, 2022
<u>AB</u>		<u>2MG</u>	<u>A216004 003</u>	Nov 18, 2022
<u>AB</u>		<u>5MG</u>	<u>A216004 004</u>	Nov 18, 2022
<u>AB</u>		<u>10MG</u>	<u>A216004 005</u>	Nov 18, 2022
<u>AB</u>		<u>20MG</u>	<u>A216004 006</u>	Nov 18, 2022
<u>AB</u>	MYLAN	<u>0.5MG</u>	<u>A070278 006</u>	Jun 10, 1986
<u>AB</u>		<u>1MG</u>	<u>A070278 004</u>	Jun 10, 1986
<u>AB</u>	!	<u>2MG</u>	<u>A070278 001</u>	Jun 10, 1986
<u>AB</u>		<u>5MG</u>	<u>A070278 005</u>	Jun 10, 1986
<u>AB</u>		<u>10MG</u>	<u>A070278 002</u>	Jul 16, 2009
<u>AB</u>		<u>20MG</u>	<u>A070278 003</u>	Jul 16, 2009
<u>AB</u>	UPSHER SMITH LABS	<u>0.5MG</u>	<u>A211061 001</u>	Jan 08, 2020
<u>AB</u>		<u>1MG</u>	<u>A211061 002</u>	Jan 08, 2020
<u>AB</u>		<u>2MG</u>	<u>A211061 003</u>	Jan 08, 2020
<u>AB</u>		<u>5MG</u>	<u>A211061 004</u>	Jan 08, 2020
<u>AB</u>		<u>10MG</u>	<u>A211061 005</u>	Jan 08, 2020
<u>AB</u>		<u>20MG</u>	<u>A211061 006</u>	Jan 08, 2020
<u>AB</u>	ZYDUS PHARMS USA	<u>0.5MG</u>	<u>A077580 001</u>	Jan 17, 2023
<u>AB</u>		<u>1MG</u>	<u>A077580 002</u>	Jan 17, 2023
<u>AB</u>		<u>2MG</u>	<u>A077580 006</u>	Jan 17, 2023
<u>AB</u>		<u>5MG</u>	<u>A077580 003</u>	Nov 29, 2007
<u>AB</u>		<u>10MG</u>	<u>A077580 004</u>	Nov 29, 2007
<u>AB</u>		<u>20MG</u>	<u>A077580 005</u>	Nov 29, 2007

HALOPERIDOL DECANOATE

INJECTABLE; INJECTION

HALDOL

<u>AO</u>	+!	JANSSEN PHARMS	<u>EQ 50MG BASE/ML</u>	<u>N018701 001</u>	Jan 14, 1986
<u>AO</u>	+!		<u>EQ 100MG BASE/ML</u>	<u>N018701 002</u>	Jan 31, 1997

HALOPERIDOL DECANOATE

<u>AO</u>	FRESENIUS KABI USA	<u>EQ 50MG BASE/ML</u>	<u>A074893 001</u>	Dec 19, 1997
<u>AO</u>		<u>EQ 100MG BASE/ML</u>	<u>A074893 002</u>	Dec 19, 1997
<u>AO</u>	GLAND PHARMA LTD	<u>EQ 50MG BASE/ML</u>	<u>A205241 001</u>	May 12, 2017
<u>AO</u>		<u>EQ 100MG BASE/ML</u>	<u>A205241 002</u>	May 12, 2017
<u>AO</u>	HIKMA	<u>EQ 50MG BASE/ML</u>	<u>A074811 001</u>	Jan 30, 1998
<u>AO</u>		<u>EQ 100MG BASE/ML</u>	<u>A075305 001</u>	Sep 28, 1998
<u>AO</u>	MANKIND PHARMA	<u>EQ 50MG BASE/ML</u>	<u>A216730 001</u>	May 23, 2023
<u>AO</u>		<u>EQ 100MG BASE/ML</u>	<u>A216730 002</u>	May 23, 2023

PRESCRIPTION DRUG PRODUCT LIST

HALOPERIDOL DECANOATE

INJECTABLE; INJECTION

HALOPERIDOL DECANOATE

<u>AO</u>	MEITHEAL	<u>EQ 50MG BASE/ML</u>	<u>A214507 001</u>	Jul 26, 2021
<u>AO</u>		<u>EQ 100MG BASE/ML</u>	<u>A214507 002</u>	Jul 26, 2021
<u>AO</u>	MYLAN LABS LTD	<u>EQ 50MG BASE/ML</u>	<u>A075440 001</u>	Feb 28, 2000
<u>AO</u>		<u>EQ 100MG BASE/ML</u>	<u>A075440 002</u>	Feb 28, 2000
<u>AO</u>	SOMERSET THERAPS LLC	<u>EQ 50MG BASE/ML</u>	<u>A209101 001</u>	Jul 03, 2018
<u>AO</u>		<u>EQ 100MG BASE/ML</u>	<u>A209101 002</u>	Jul 03, 2018
<u>AO</u>	ZYDUS PHARMS	<u>EQ 50MG BASE/ML</u>	<u>A211180 001</u>	Oct 22, 2019
<u>AO</u>		<u>EQ 100MG BASE/ML</u>	<u>A211180 002</u>	Oct 22, 2019

HALOPERIDOL LACTATE

CONCENTRATE; ORAL

HALOPERIDOL

<u>AA</u>	LANNETT CO INC	<u>EQ 2MG BASE/ML</u>	<u>A073364 001</u>	Sep 28, 1993
<u>AA</u>	! PHARM ASSOC	<u>EQ 2MG BASE/ML</u>	<u>A073037 001</u>	Feb 26, 1993
<u>AA</u>	RUBICON	<u>EQ 2MG BASE/ML</u>	<u>A218371 001</u>	Jan 31, 2024

INJECTABLE; INJECTION

HALOPERIDOL

<u>AP</u>	FRESENIUS KABI USA	<u>EQ 5MG BASE/ML</u>	<u>A075689 001</u>	Mar 09, 2001
<u>AP</u>		<u>EQ 5MG BASE/ML</u>	<u>A210356 001</u>	Jul 01, 2019
<u>AP</u>	GLAND PHARMA LTD	<u>EQ 5MG BASE/ML</u>	<u>A076774 001</u>	Aug 25, 2004
<u>AP</u>	HIKMA	<u>EQ 5MG BASE/ML</u>	<u>A075858 001</u>	Jun 18, 2001
<u>AP</u>	! MYLAN LABS LTD	<u>EQ 5MG BASE/ML</u>	<u>A078347 001</u>	Sep 14, 2009
<u>AP</u>	SAGENT PHARMS	<u>EQ 5MG BASE/ML</u>	<u>A091637 001</u>	Sep 02, 2011
<u>AP</u>		<u>EQ 5MG BASE/ML</u>	<u>A200742 001</u>	Sep 02, 2011

HEPARIN SODIUM

INJECTABLE; INJECTION

HEPARIN SODIUM

<u>AP</u>	B BRAUN MEDICAL INC	<u>5,000 UNITS/0.5ML</u>	<u>A208827 001</u>	Nov 19, 2018
<u>AP</u>	BE PHARMS	<u>1,000 UNITS/ML</u>	<u>A214804 001</u>	Dec 29, 2020
<u>AP</u>		<u>10,000 UNITS/ML</u>	<u>A214839 001</u>	Dec 29, 2020
<u>AP</u>	+! FRESENIUS KABI USA	<u>1,000 UNITS/ML</u>	<u>N017029 001</u>	
<u>AP</u>		<u>5,000 UNITS/ML</u>	<u>A206552 001</u>	Jun 10, 2016
<u>AP</u>	+!	<u>5,000 UNITS/ML</u>	<u>N017651 006</u>	
<u>AP</u>	+!	<u>10,000 UNITS/ML</u>	<u>N017029 003</u>	
<u>AP</u>	+!	<u>20,000 UNITS/ML</u>	<u>N017029 004</u>	
<u>AP</u>	GLAND	<u>1,000 UNITS/ML</u>	<u>A205323 002</u>	Nov 18, 2019
<u>AP</u>		<u>5,000 UNITS/ML</u>	<u>A205323 001</u>	Feb 06, 2017
<u>AP</u>	+! HIKMA	<u>1,000 UNITS/ML</u>	<u>N017037 001</u>	
<u>AP</u>	+!	<u>5,000 UNITS/ML</u>	<u>N017037 002</u>	
<u>AP</u>		<u>5,000 UNITS/0.5ML</u>	<u>N017037 013</u>	Apr 07, 1986
<u>AP</u>	+!	<u>10,000 UNITS/ML</u>	<u>N017037 003</u>	
<u>AP</u>	HOSPIRA INC	<u>1,000 UNITS/ML</u>	<u>A090571 001</u>	Aug 31, 2009
<u>AP</u>		<u>5,000 UNITS/ML</u>	<u>A090571 002</u>	Aug 31, 2009
<u>AP</u>		<u>10,000 UNITS/ML</u>	<u>A090571 003</u>	Aug 31, 2009
<u>AP</u>	MYLAN LABS LTD	<u>1,000 UNITS/ML</u>	<u>A203851 001</u>	Nov 30, 2017
<u>AP</u>		<u>5,000 UNITS/ML</u>	<u>A203851 002</u>	Nov 30, 2017
<u>AP</u>		<u>10,000 UNITS/ML</u>	<u>A203851 003</u>	Nov 30, 2017
<u>AP</u>		<u>20,000 UNITS/ML</u>	<u>A203852 001</u>	Nov 30, 2017
<u>AP</u>	NANJING KING-FRIEND	<u>1,000 UNITS/ML</u>	<u>A211005 001</u>	Dec 14, 2018
<u>AP</u>		<u>1,000 UNITS/ML</u>	<u>A211007 001</u>	May 28, 2019
<u>AP</u>		<u>5,000 UNITS/ML</u>	<u>A211007 002</u>	May 28, 2019
<u>AP</u>		<u>5,000 UNITS/ML</u>	<u>A212061 001</u>	Jul 15, 2020
<u>AP</u>		<u>10,000 UNITS/ML</u>	<u>A211007 003</u>	May 28, 2019
<u>AP</u>		<u>20,000 UNITS/ML</u>	<u>A211004 001</u>	Feb 24, 2020
<u>AP</u>	SAGENT PHARMS	<u>1,000 UNITS/ML</u>	<u>A090808 001</u>	Jun 30, 2010
<u>AP</u>		<u>5,000 UNITS/ML</u>	<u>A090808 002</u>	Jun 30, 2010
<u>AP</u>		<u>10,000 UNITS/ML</u>	<u>A090808 003</u>	Jun 30, 2010
<u>AP</u>		<u>20,000 UNITS/ML</u>	<u>A090809 001</u>	Jun 30, 2010
<u>AP</u>	SANDOZ	<u>1,000 UNITS/ML</u>	<u>A091682 001</u>	Jun 08, 2011
<u>AP</u>		<u>5,000 UNITS/ML</u>	<u>A091682 002</u>	Jun 08, 2011
<u>AP</u>		<u>10,000 UNITS/ML</u>	<u>A201002 001</u>	Jun 08, 2011
<u>AP</u>	SHENZHEN TECHDOW	<u>1,000 UNITS/ML</u>	<u>A202957 001</u>	Jun 12, 2014
<u>AP</u>		<u>5,000 UNITS/ML</u>	<u>A202733 001</u>	Jun 12, 2014
<u>AP</u>		<u>5,000 UNITS/ML</u>	<u>A202957 002</u>	Jun 12, 2014
<u>AP</u>		<u>10,000 UNITS/ML</u>	<u>A203198 001</u>	Jun 12, 2014
<u>AP</u>		<u>20,000 UNITS/ML</u>	<u>A203198 002</u>	Jun 12, 2014

HEPARIN SODIUM 1,000 UNITS AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

<u>AP</u>	BAXTER HLTHCARE	<u>200 UNITS/100ML</u>	<u>N018609 001</u>	Apr 28, 1982
-----------	-----------------	------------------------	--------------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

HEPARIN SODIUM

INJECTABLE; INJECTION

HEPARIN SODIUM 1,000 UNITS IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

<u>AP</u>	<u>+!</u>	<u>B BRAUN</u>	<u>200 UNITS/100ML</u>	<u>N019953</u>	<u>001</u>	Jul 20, 1992
<u>AP</u>		<u>FRESENIUS KABI USA</u>	<u>200 UNITS/100ML</u>	<u>A212441</u>	<u>001</u>	Jul 24, 2020
<u>AP</u>	<u>+!</u>	<u>HOSPIRA</u>	<u>200 UNITS/100ML</u>	<u>N018916</u>	<u>010</u>	Jun 23, 1989

HEPARIN SODIUM 10,000 UNITS IN DEXTROSE 5% IN PLASTIC CONTAINER

<u>AP</u>		<u>HOSPIRA</u>	<u>10,000 UNITS/100ML</u>	<u>N019339</u>	<u>003</u>	Mar 27, 1985
-----------	--	----------------	---------------------------	----------------	------------	--------------

HEPARIN SODIUM 2,000 UNITS AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

<u>AP</u>		<u>BAXTER HLTHCARE</u>	<u>200 UNITS/100ML</u>	<u>N018609</u>	<u>002</u>	Apr 28, 1982
-----------	--	------------------------	------------------------	----------------	------------	--------------

HEPARIN SODIUM 2,000 UNITS IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

<u>AP</u>		<u>FRESENIUS KABI USA</u>	<u>200 UNITS/100ML</u>	<u>A212441</u>	<u>002</u>	Jul 24, 2020
<u>AP</u>	<u>+!</u>	<u>HOSPIRA</u>	<u>200 UNITS/100ML</u>	<u>N018916</u>	<u>011</u>	Jun 23, 1989

HEPARIN SODIUM 25,000 UNITS IN DEXTROSE 5% IN PLASTIC CONTAINER

<u>AP</u>	<u>+!</u>	<u>B BRAUN</u>	<u>5,000 UNITS/100ML</u>	<u>N019952</u>	<u>004</u>	Jul 20, 1992
<u>AP</u>	<u>+!</u>		<u>10,000 UNITS/100ML</u>	<u>N019952</u>	<u>005</u>	Jul 20, 1992
<u>AP</u>		<u>HOSPIRA</u>	<u>5,000 UNITS/100ML</u>	<u>N019339</u>	<u>004</u>	Mar 27, 1985
<u>AP</u>			<u>10,000 UNITS/100ML</u>	<u>N019339</u>	<u>002</u>	Mar 27, 1985

HEPARIN SODIUM IN PLASTIC CONTAINER

<u>AP</u>	<u>+!</u>	<u>FRESENIUS KABI USA</u>	<u>1,000 UNITS/ML</u>	<u>N017029</u>	<u>013</u>	Dec 05, 1985
<u>AP</u>	<u>+!</u>		<u>5,000 UNITS/ML</u>	<u>N017029</u>	<u>014</u>	Dec 05, 1985
<u>AP</u>	<u>+!</u>		<u>10,000 UNITS/ML</u>	<u>N017029</u>	<u>015</u>	Dec 05, 1985
<u>AP</u>	<u>+!</u>		<u>20,000 UNITS/ML</u>	<u>N017029</u>	<u>016</u>	Dec 05, 1985

HEPARIN SODIUM PRESERVATIVE FREE

<u>AP</u>	<u>+!</u>	<u>FRESENIUS KABI USA</u>	<u>1,000 UNITS/ML</u>	<u>N017029</u>	<u>010</u>	Apr 28, 1986
<u>AP</u>	<u>+!</u>		<u>10,000 UNITS/ML</u>	<u>N017029</u>	<u>019</u>	Nov 22, 2010
<u>AP</u>		<u>HOSPIRA</u>	<u>10,000 UNITS/ML</u>	<u>A089522</u>	<u>001</u>	May 04, 1987
<u>AP</u>		<u>NANJING KING-FRIEND</u>	<u>10,000 UNITS/ML</u>	<u>A212060</u>	<u>001</u>	Apr 02, 2020
<u>AP</u>		<u>SAGENT PHARMS</u>	<u>1,000 UNITS/ML</u>	<u>A090810</u>	<u>001</u>	Jun 30, 2010
<u>AP</u>		<u>SHENZHEN TECHDOW</u>	<u>1,000 UNITS/ML</u>	<u>A202732</u>	<u>001</u>	Jun 12, 2014

HEPARIN SODIUM

	<u>+!</u>	<u>FRESENIUS KABI USA</u>	<u>10,000 UNITS/ML</u>	<u>N017029</u>	<u>020</u>	Mar 31, 2011
	<u>!</u>	<u>HOSPIRA</u>	<u>5,000 UNITS/ML</u>	<u>A088100</u>	<u>001</u>	Apr 28, 1983
	<u>+!</u>	<u>PFIZER</u>	<u>1,000 UNITS/ML</u>	<u>N201370</u>	<u>001</u>	Jul 21, 2011
	<u>+!</u>		<u>5,000 UNITS/ML</u>	<u>N201370</u>	<u>002</u>	Jul 21, 2011

HEPARIN SODIUM 12,500 UNITS IN DEXTROSE 5% IN PLASTIC CONTAINER

		<u>HOSPIRA</u>	<u>5,000 UNITS/100ML</u>	<u>N019339</u>	<u>001</u>	Mar 27, 1985
--	--	----------------	--------------------------	----------------	------------	--------------

HEPARIN SODIUM 12,500 UNITS IN SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER

	<u>+!</u>	<u>HOSPIRA</u>	<u>5,000 UNITS/100ML</u>	<u>N018916</u>	<u>006</u>	Jan 31, 1984
--	-----------	----------------	--------------------------	----------------	------------	--------------

HEPARIN SODIUM 2,000 UNITS IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

	<u>+!</u>	<u>B BRAUN</u>	<u>200 UNITS/100ML</u>	<u>N019953</u>	<u>002</u>	Dec 15, 2023
--	-----------	----------------	------------------------	----------------	------------	--------------

HEPARIN SODIUM 20,000 UNITS IN DEXTROSE 5% IN PLASTIC CONTAINER

	<u>+!</u>	<u>B BRAUN</u>	<u>4,000 UNITS/100ML</u>	<u>N019952</u>	<u>001</u>	Jul 20, 1992
--	-----------	----------------	--------------------------	----------------	------------	--------------

HEPARIN SODIUM 25,000 UNITS IN DEXTROSE 5%

	<u>+!</u>	<u>FRESENIUS KABI USA</u>	<u>50 UNITS/ML</u>	<u>N017029</u>	<u>022</u>	Aug 24, 2017
	<u>+!</u>		<u>100 UNITS/ML</u>	<u>N017029</u>	<u>023</u>	Aug 24, 2017

HEPARIN SODIUM 25,000 UNITS IN SODIUM CHLORIDE 0.45%

	<u>+!</u>	<u>FRESENIUS KABI USA</u>	<u>50 UNITS/ML</u>	<u>N017029</u>	<u>024</u>	Aug 24, 2017
	<u>+!</u>		<u>100 UNITS/ML</u>	<u>N017029</u>	<u>025</u>	Aug 24, 2017

HEPARIN SODIUM 25,000 UNITS IN SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER

	<u>+!</u>	<u>HOSPIRA</u>	<u>5,000 UNITS/100ML</u>	<u>N018916</u>	<u>007</u>	Jan 31, 1984
	<u>+!</u>		<u>10,000 UNITS/100ML</u>	<u>N018916</u>	<u>008</u>	Jan 31, 1984

HEPARIN SODIUM PRESERVATIVE FREE

	<u>+!</u>	<u>PFIZER</u>	<u>1,000 UNITS/ML</u>	<u>N201370</u>	<u>004</u>	Jul 21, 2011
--	-----------	---------------	-----------------------	----------------	------------	--------------

HEPARIN SODIUM; TAUROLIDINE

SOLUTION; N/A

DEFENCATH

	<u>+!</u>	<u>CORMEDIX</u>	<u>3,000 UNITS/3ML (1,000 UNITS/ML); 40.5MG/3ML (13.5MG/ML)</u>	<u>N214520</u>	<u>001</u>	Nov 15, 2023
	<u>+!</u>		<u>5,000 UNITS/5ML (1,000 UNITS/ML); 67.5MG/5ML (13.5MG/ML)</u>	<u>N214520</u>	<u>002</u>	Nov 15, 2023

HEXACHLOROPHENE

SPONGE; TOPICAL

PRE-OP

<u>AT</u>	<u>+!</u>	<u>DAVIS AND GECK</u>	<u>480MG</u>	<u>N017433</u>	<u>001</u>	
-----------	-----------	-----------------------	--------------	----------------	------------	--

PRE-OP II

<u>AT</u>	<u>+</u>	<u>DAVIS AND GECK</u>	<u>480MG</u>	<u>N017433</u>	<u>002</u>	
-----------	----------	-----------------------	--------------	----------------	------------	--

PRESCRIPTION DRUG PRODUCT LISTHEXAMINOLEVULINATE HYDROCHLORIDEFOR SOLUTION; INTRAVESICAL
CYSVIEW KIT

+! PHOTOCURE ASA 100MG/VIAL N022555 001 May 28, 2010

HISTRELIN ACETATEIMPLANT; SUBCUTANEOUS
SUPPRELIN LA

+! ENDO OPERATIONS 50MG N022058 001 May 03, 2007

HOMATROPINE METHYLBROMIDE; HYDROCODONE BITARTRATE

SYRUP; ORAL

HYCODAN**AA** + GENUS 1.5MG/5ML; 5MG/5ML **N005213 002** Jul 26, 1988HYDROCODONE BITARTRATE AND HOMATROPINE METHYLBROMIDE**AA** ! ABHAI LLC 1.5MG/5ML; 5MG/5ML **A207487 001** Feb 21, 2017**AA** ACTAVIS MID 1.5MG/5ML; 5MG/5ML **A088017 001** Jul 05, 1983

ATLANTIC

AA NOVEL LABS INC 1.5MG/5ML; 5MG/5ML **A203535 001** Feb 13, 2017**AA** PADAGIS US 1.5MG/5ML; 5MG/5ML **A205731 001** Feb 15, 2017**AA** WOCKHARDT BIO AG 1.5MG/5ML; 5MG/5ML **A088008 001** Mar 03, 1983

TABLET; ORAL

HOMATROPINE METHYLBROMIDE AND HYDROCODONE BITARTRATE**AA** ! AVANTHI INC 1.5MG; 5MG **A207176 001** Aug 07, 2017HYCODAN**AA** + GENUS 1.5MG; 5MG **N005213 001** Jul 26, 1988HYDRALAZINE HYDROCHLORIDE

INJECTABLE; INJECTION

HYDRALAZINE HYDROCHLORIDE**AP** ADRASTEIA PHARMA 20MG/ML **A203110 001** Jun 29, 2015**AP** AM REGENT 20MG/ML **A040136 001** Jun 30, 1997**AP** EUGIA PHARMA 20MG/ML **A215147 001** Feb 13, 2023**AP** ! FRESENIUS KABI USA 20MG/ML **A040388 001** Mar 13, 2001**AP** HIKMA 20MG/ML **A213667 001** Dec 18, 2020**AP** NAVINTA LLC 20MG/ML **A202938 001** Mar 28, 2013**AP** RISING 20MG/ML **A040730 001** Apr 21, 2009

TABLET; ORAL

HYDRALAZINE HYDROCHLORIDE**AA** ALKEM LABS LTD 10MG **A200737 001** Dec 07, 2012**AA** 25MG **A200737 002** Dec 07, 2012**AA** 50MG **A200737 003** Dec 07, 2012**AA** 100MG **A200737 004** Dec 07, 2012**AA** CADILA PHARMS LTD 25MG **A203845 001** Sep 18, 2014**AA** 50MG **A203845 002** Sep 18, 2014**AA** 100MG **A203845 003** Sep 18, 2014**AA** GLENMARK PHARMS LTD 10MG **A090527 001** May 29, 2009**AA** 25MG **A090527 002** May 29, 2009**AA** 50MG **A090527 003** May 29, 2009**AA** 100MG **A090527 004** May 29, 2009**AA** HERITAGE 10MG **A086242 001** Feb 04, 2010**AA** 25MG **A086242 003****AA** 50MG **A086242 002****AA** 100MG **A086242 004** Feb 04, 2010**AA** HERITAGE PHARMS INC 10MG **A040858 001** Feb 26, 2010**AA** 25MG **A040858 002** Feb 26, 2010**AA** 50MG **A040858 003** Feb 26, 2010**AA** 100MG **A040858 004** Feb 26, 2010**AA** HETERO LABS LTD III 10MG **A040901 001** Sep 12, 2008**AA** 25MG **A040901 002** Sep 12, 2008**AA** 50MG **A040901 003** Sep 12, 2008**AA** 100MG **A040901 004** Sep 12, 2008**AA** MACLEODS PHARMS LTD 10MG **A211010 001** Aug 21, 2024**AA** 25MG **A211010 002** Aug 21, 2024**AA** 50MG **A211010 003** Aug 21, 2024**AA** 100MG **A211010 004** Aug 21, 2024**AA** ! PLIVA 10MG **A089097 001** Dec 18, 1985**AA** +! 25MG **A088467 001** May 01, 1984**AA** +! 50MG **A088468 001** May 01, 1984**AA** ! 100MG **A089098 001** Dec 18, 1985**AA** SCIEGEN PHARMS INC 10MG **A205236 001** May 26, 2017**AA** 25MG **A205236 002** May 26, 2017**AA** 50MG **A205236 003** May 26, 2017**AA** 100MG **A205236 004** May 26, 2017

PRESCRIPTION DRUG PRODUCT LIST

HYDRALAZINE HYDROCHLORIDE

TABLET; ORAL

HYDRALAZINE HYDROCHLORIDE

<u>AA</u>	STRIDES PHARMA	<u>10MG</u>	<u>A087836</u>	<u>001</u>	Oct 05, 1982
<u>AA</u>		<u>25MG</u>	<u>A086961</u>	<u>002</u>	
<u>AA</u>		<u>25MG</u>	<u>A200770</u>	<u>001</u>	May 03, 2013
<u>AA</u>		<u>50MG</u>	<u>A086962</u>	<u>001</u>	
<u>AA</u>		<u>50MG</u>	<u>A200770</u>	<u>002</u>	May 03, 2013
<u>AA</u>		<u>100MG</u>	<u>A088391</u>	<u>001</u>	Sep 27, 1983
<u>AA</u>		<u>100MG</u>	<u>A200770</u>	<u>003</u>	May 03, 2013

HYDRALAZINE HYDROCHLORIDE; HYDROCHLOROTHIAZIDE

CAPSULE; ORAL

HYDRA-ZIDE

	STRIDES PHARMA	25MG; 25MG	A088957	001	Oct 21, 1985
!		50MG; 50MG	A088946	001	Oct 21, 1985

HYDRALAZINE HYDROCHLORIDE; ISOSORBIDE DINITRATE

TABLET; ORAL

BIDIL

<u>AB</u>	+!	AZURITY	<u>37.5MG; 20MG</u>	<u>N020727</u>	<u>001</u>	Jun 23, 2005
-----------	----	---------	---------------------	----------------	------------	--------------

ISOSORBIDE DINITRATE AND HYDRALAZINE HYDROCHLORIDE

<u>AB</u>		I3 PHARMS	<u>37.5MG; 20MG</u>	<u>A215988</u>	<u>001</u>	Jan 17, 2024
<u>AB</u>		RICONPHARMA LLC	<u>37.5MG; 20MG</u>	<u>A215586</u>	<u>001</u>	Apr 06, 2022

HYDROCHLOROTHIAZIDE

CAPSULE; ORAL

HYDROCHLOROTHIAZIDE

<u>AB</u>		AUROBINDO PHARMA	<u>12.5MG</u>	<u>A078164</u>	<u>001</u>	Sep 18, 2007
<u>AB</u>		JUBILANT CADISTA	<u>12.5MG</u>	<u>A078391</u>	<u>001</u>	Feb 11, 2008
<u>AB</u>		PRINSTON INC	<u>12.5MG</u>	<u>A075907</u>	<u>001</u>	Sep 17, 2002
<u>AB</u>		SCIEGEN PHARMS INC	<u>12.5MG</u>	<u>A203561</u>	<u>001</u>	Jan 14, 2019
<u>AB</u>		UNICHEM	<u>12.5MG</u>	<u>A090510</u>	<u>001</u>	Jan 19, 2010

MICROZIDE

<u>AB</u>	+!	TEVA BRANDED PHARM	<u>12.5MG</u>	<u>N020504</u>	<u>001</u>	Dec 27, 1996
-----------	----	--------------------	---------------	----------------	------------	--------------

TABLET; ORAL

HYDROCHLOROTHIAZIDE

<u>AB</u>		ACCORD HLTHCARE	<u>12.5MG</u>	<u>A202556</u>	<u>001</u>	Sep 24, 2012
<u>AB</u>			<u>25MG</u>	<u>A202556</u>	<u>002</u>	Sep 24, 2012
<u>AB</u>			<u>50MG</u>	<u>A202556</u>	<u>003</u>	Sep 24, 2012
<u>AB</u>		ACTAVIS ELIZABETH	<u>12.5MG</u>	<u>A040707</u>	<u>001</u>	Feb 27, 2007
<u>AB</u>		AUROBINDO PHARMA	<u>25MG</u>	<u>A040780</u>	<u>001</u>	Jul 20, 2007
<u>AB</u>			<u>50MG</u>	<u>A040780</u>	<u>002</u>	Jul 20, 2007
<u>AB</u>		HERITAGE	<u>12.5MG</u>	<u>A085182</u>	<u>003</u>	May 02, 2023
<u>AB</u>			<u>25MG</u>	<u>A085182</u>	<u>002</u>	
<u>AB</u>			<u>50MG</u>	<u>A085182</u>	<u>001</u>	
<u>AB</u>	+	IVAX SUB TEVA PHARMS	<u>12.5MG</u>	<u>A083177</u>	<u>003</u>	Sep 19, 2024
<u>AB</u>	+		<u>25MG</u>	<u>A083177</u>	<u>001</u>	
<u>AB</u>	+!		<u>50MG</u>	<u>A083177</u>	<u>002</u>	
<u>AB</u>		LEADING	<u>12.5MG</u>	<u>A040702</u>	<u>003</u>	May 10, 2017
<u>AB</u>			<u>25MG</u>	<u>A040702</u>	<u>001</u>	Mar 16, 2007
<u>AB</u>			<u>50MG</u>	<u>A040702</u>	<u>002</u>	Mar 16, 2007
<u>AB</u>		OXFORD PHARMS	<u>25MG</u>	<u>A087059</u>	<u>001</u>	
<u>AB</u>			<u>50MG</u>	<u>A087068</u>	<u>001</u>	
<u>AB</u>		SCIEGEN PHARMS INC	<u>25MG</u>	<u>A203018</u>	<u>001</u>	Jul 23, 2014
<u>AB</u>			<u>50MG</u>	<u>A203018</u>	<u>002</u>	Jul 23, 2014
<u>AB</u>		UNICHEM	<u>25MG</u>	<u>A040907</u>	<u>001</u>	Aug 15, 2008
<u>AB</u>			<u>50MG</u>	<u>A040907</u>	<u>002</u>	Aug 15, 2008

HYDROCHLOROTHIAZIDE; IRBESARTAN

TABLET; ORAL

AVALIDE

<u>AB</u>	+!	SANOVI AVENTIS US	<u>12.5MG; 150MG</u>	<u>N020758</u>	<u>002</u>	Sep 30, 1997
<u>AB</u>	+!		<u>12.5MG; 300MG</u>	<u>N020758</u>	<u>003</u>	Aug 31, 1998

IRBESARTAN AND HYDROCHLOROTHIAZIDE

<u>AB</u>		ALEMBIC	<u>12.5MG; 150MG</u>	<u>A091370</u>	<u>001</u>	Oct 15, 2012
<u>AB</u>			<u>12.5MG; 300MG</u>	<u>A091370</u>	<u>002</u>	Oct 15, 2012
<u>AB</u>			<u>25MG; 300MG</u>	<u>A091370</u>	<u>003</u>	Oct 12, 2016
<u>AB</u>		AUROBINDO PHARMA LTD	<u>12.5MG; 150MG</u>	<u>A203630</u>	<u>001</u>	Feb 22, 2013
<u>AB</u>			<u>12.5MG; 300MG</u>	<u>A203630</u>	<u>002</u>	Feb 22, 2013
<u>AB</u>			<u>25MG; 300MG</u>	<u>A203630</u>	<u>003</u>	Mar 31, 2016
<u>AB</u>		DR REDDYS LABS LTD	<u>12.5MG; 150MG</u>	<u>A203500</u>	<u>001</u>	Sep 27, 2012
<u>AB</u>			<u>12.5MG; 300MG</u>	<u>A203500</u>	<u>002</u>	Sep 27, 2012

PRESCRIPTION DRUG PRODUCT LIST

HYDROCHLOROTHIAZIDE; IRBESARTAN

TABLET; ORAL

IRBESARTAN AND HYDROCHLOROTHIAZIDE

<u>AB</u>	HIKMA	<u>12.5MG;150MG</u>	<u>A090351 001</u>	Oct 15, 2012
<u>AB</u>		<u>12.5MG;300MG</u>	<u>A090351 002</u>	Oct 15, 2012
<u>AB</u>		<u>25MG;300MG</u>	<u>A090351 003</u>	Jun 08, 2017
<u>AB</u>	HISUN PHARM HANGZHOU	<u>12.5MG;150MG</u>	<u>A207896 001</u>	Oct 14, 2016
<u>AB</u>		<u>12.5MG;300MG</u>	<u>A207896 002</u>	Oct 14, 2016
<u>AB</u>	LUPIN LTD	<u>12.5MG;150MG</u>	<u>A201524 001</u>	Feb 27, 2013
<u>AB</u>		<u>12.5MG;300MG</u>	<u>A201524 002</u>	Feb 27, 2013
<u>AB</u>	MACLEODS PHARMS LTD	<u>12.5MG;150MG</u>	<u>A202414 001</u>	Sep 27, 2012
<u>AB</u>		<u>12.5MG;300MG</u>	<u>A202414 002</u>	Sep 27, 2012
<u>AB</u>	PRINSTON INC	<u>12.5MG;150MG</u>	<u>A203072 001</u>	May 09, 2014
<u>AB</u>		<u>12.5MG;300MG</u>	<u>A203072 002</u>	May 09, 2014
<u>AB</u>	SANDOZ	<u>12.5MG;150MG</u>	<u>A077446 001</u>	Sep 27, 2012
<u>AB</u>		<u>12.5MG;300MG</u>	<u>A077446 002</u>	Sep 27, 2012
<u>AB</u>	TEVA	<u>12.5MG;150MG</u>	<u>A077369 001</u>	Mar 30, 2012
<u>AB</u>		<u>12.5MG;300MG</u>	<u>A077369 002</u>	Mar 30, 2012

HYDROCHLOROTHIAZIDE; LISINAPRIL

TABLET; ORAL

LISINAPRIL AND HYDROCHLOROTHIAZIDE

<u>AB</u>	AUROBINDO	<u>12.5MG;10MG</u>	<u>A077606 001</u>	Mar 14, 2006
<u>AB</u>		<u>12.5MG;20MG</u>	<u>A077606 002</u>	Mar 14, 2006
<u>AB</u>		<u>25MG;20MG</u>	<u>A077606 003</u>	Mar 14, 2006
<u>AB</u>	COREPHARMA	<u>12.5MG;10MG</u>	<u>A076674 001</u>	Oct 05, 2004
<u>AB</u>		<u>12.5MG;20MG</u>	<u>A076674 002</u>	Oct 05, 2004
<u>AB</u>		<u>25MG;20MG</u>	<u>A076674 003</u>	Oct 05, 2004
<u>AB</u>	INVAGEN PHARMS	<u>12.5MG;10MG</u>	<u>A204058 001</u>	May 23, 2017
<u>AB</u>	!	<u>12.5MG;20MG</u>	<u>A204058 002</u>	May 23, 2017
<u>AB</u>	!	<u>25MG;20MG</u>	<u>A204058 003</u>	May 23, 2017
<u>AB</u>	LUPIN	<u>12.5MG;10MG</u>	<u>A077912 001</u>	Sep 27, 2006
<u>AB</u>		<u>12.5MG;20MG</u>	<u>A077912 002</u>	Sep 27, 2006
<u>AB</u>		<u>25MG;20MG</u>	<u>A077912 003</u>	Sep 27, 2006
<u>AB</u>	PRINSTON INC	<u>12.5MG;10MG</u>	<u>A076230 001</u>	Jul 01, 2002
<u>AB</u>		<u>12.5MG;20MG</u>	<u>A076230 002</u>	Jul 01, 2002
<u>AB</u>		<u>25MG;20MG</u>	<u>A076230 003</u>	Jul 01, 2002
<u>AB</u>	SANDOZ	<u>12.5MG;10MG</u>	<u>A076262 001</u>	Jul 01, 2002
<u>AB</u>		<u>12.5MG;20MG</u>	<u>A076262 002</u>	Jul 01, 2002
<u>AB</u>		<u>25MG;20MG</u>	<u>A076262 003</u>	Jul 01, 2002
<u>AB</u>	SUN PHARM INDS LTD	<u>12.5MG;10MG</u>	<u>A076007 001</u>	Jul 01, 2002
<u>AB</u>		<u>12.5MG;20MG</u>	<u>A076007 002</u>	Jul 01, 2002
<u>AB</u>		<u>25MG;20MG</u>	<u>A076007 003</u>	Jul 01, 2002
<u>AB</u>	WATSON LABS	<u>12.5MG;10MG</u>	<u>A076194 003</u>	Jul 01, 2002
<u>AB</u>		<u>12.5MG;20MG</u>	<u>A076194 001</u>	Jul 01, 2002
<u>AB</u>		<u>25MG;20MG</u>	<u>A076194 002</u>	Jul 01, 2002
<u>ZESTORETIC</u>				
<u>AB</u>	+ ALMATICA	<u>12.5MG;10MG</u>	<u>N019888 003</u>	Nov 18, 1993
<u>AB</u>	+	<u>12.5MG;20MG</u>	<u>N019888 001</u>	Sep 20, 1990
<u>AB</u>	+	<u>25MG;20MG</u>	<u>N019888 002</u>	Jul 20, 1989

HYDROCHLOROTHIAZIDE; LOSARTAN POTASSIUM

TABLET; ORAL

HYZAAR

<u>AB</u>	+ ORGANON	<u>12.5MG;50MG</u>	<u>N020387 001</u>	Apr 28, 1995
<u>AB</u>	+	<u>12.5MG;100MG</u>	<u>N020387 003</u>	Oct 20, 2005
<u>AB</u>	+	<u>25MG;100MG</u>	<u>N020387 002</u>	Nov 10, 1998

LOSARTAN POTASSIUM AND HYDROCHLOROTHIAZIDE

<u>AB</u>	ALEMBIC PHARMS LTD	<u>12.5MG;50MG</u>	<u>A091617 001</u>	Feb 17, 2012
<u>AB</u>		<u>12.5MG;100MG</u>	<u>A091617 002</u>	Feb 17, 2012
<u>AB</u>		<u>25MG;100MG</u>	<u>A091617 003</u>	Feb 17, 2012
<u>AB</u>	AUROBINDO PHARMA	<u>12.5MG;50MG</u>	<u>A091629 001</u>	Oct 06, 2010
<u>AB</u>		<u>12.5MG;100MG</u>	<u>A091629 002</u>	Oct 06, 2010
<u>AB</u>	!	<u>25MG;100MG</u>	<u>A091629 003</u>	Jan 06, 2010
<u>AB</u>	CHARTWELL RX	<u>12.5MG;50MG</u>	<u>A077948 001</u>	Oct 06, 2010
<u>AB</u>		<u>12.5MG;100MG</u>	<u>A077948 003</u>	Aug 19, 2010
<u>AB</u>		<u>25MG;100MG</u>	<u>A077948 002</u>	Oct 06, 2010
<u>AB</u>	JUBILANT CADISTA	<u>12.5MG;50MG</u>	<u>A201845 001</u>	Sep 18, 2012
<u>AB</u>		<u>12.5MG;100MG</u>	<u>A201845 002</u>	Sep 18, 2012
<u>AB</u>		<u>25MG;100MG</u>	<u>A201845 003</u>	Sep 18, 2012
<u>AB</u>	LUPIN LTD	<u>12.5MG;50MG</u>	<u>A078245 001</u>	Oct 06, 2010
<u>AB</u>		<u>12.5MG;100MG</u>	<u>A078245 002</u>	May 21, 2010
<u>AB</u>		<u>25MG;100MG</u>	<u>A078245 003</u>	Oct 06, 2010

PRESCRIPTION DRUG PRODUCT LIST

HYDROCHLOROTHIAZIDE; LOSARTAN POTASSIUM

TABLET; ORAL

LOSARTAN POTASSIUM AND HYDROCHLOROTHIAZIDE

<u>AB</u>	MACLEODS PHARMS LTD	<u>12.5MG; 50MG</u>	<u>A202289 001</u>	Aug 09, 2012
<u>AB</u>		<u>12.5MG; 100MG</u>	<u>A202289 002</u>	Aug 09, 2012
<u>AB</u>		<u>25MG; 100MG</u>	<u>A202289 003</u>	Aug 09, 2012
<u>AB</u>	PRINSTON INC	<u>12.5MG; 50MG</u>	<u>A204901 001</u>	Nov 06, 2017
<u>AB</u>		<u>12.5MG; 100MG</u>	<u>A204901 002</u>	Nov 06, 2017
<u>AB</u>		<u>25MG; 100MG</u>	<u>A204901 003</u>	Nov 06, 2017
<u>AB</u>	TEVA PHARMS	<u>12.5MG; 50MG</u>	<u>A077157 001</u>	Apr 06, 2010
<u>AB</u>		<u>12.5MG; 100MG</u>	<u>A077157 002</u>	Apr 06, 2010
<u>AB</u>		<u>25MG; 100MG</u>	<u>A077157 003</u>	Apr 06, 2010
<u>AB</u>	UNICHEM	<u>12.5MG; 50MG</u>	<u>A204832 001</u>	Jul 21, 2017
<u>AB</u>		<u>12.5MG; 100MG</u>	<u>A204832 002</u>	Jul 21, 2017
<u>AB</u>		<u>25MG; 100MG</u>	<u>A204832 003</u>	Jul 21, 2017
<u>AB</u>	ZYDUS PHARMS USA INC	<u>12.5MG; 50MG</u>	<u>A078385 001</u>	Oct 06, 2010
<u>AB</u>		<u>25MG; 100MG</u>	<u>A078385 002</u>	Oct 06, 2010

HYDROCHLOROTHIAZIDE; METOPROLOL TARTRATE

TABLET; ORAL

LOPRESSOR HCT

<u>AB</u>	+ VALIDUS PHARMS	<u>25MG; 50MG</u>	<u>N018303 001</u>	Dec 31, 1984
-----------	------------------	-------------------	--------------------	--------------

METOPROLOL TARTRATE AND HYDROCHLOROTHIAZIDE

<u>AB</u>	ALEMBIC	<u>25MG; 50MG</u>	<u>A202870 001</u>	Nov 06, 2013
<u>AB</u>	!	<u>25MG; 100MG</u>	<u>A202870 002</u>	Nov 06, 2013
<u>AB</u>		<u>50MG; 100MG</u>	<u>A202870 003</u>	Nov 06, 2013
<u>AB</u>	MYLAN	<u>25MG; 50MG</u>	<u>A076792 001</u>	Aug 20, 2004
<u>AB</u>		<u>25MG; 100MG</u>	<u>A076792 002</u>	Aug 20, 2004
<u>AB</u>		<u>50MG; 100MG</u>	<u>A076792 003</u>	Aug 20, 2004

HYDROCHLOROTHIAZIDE; MOEXIPRIL HYDROCHLORIDE

TABLET; ORAL

MOEXIPRIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE

<u>AB</u>	GLENMARK PHARMS	<u>12.5MG; 7.5MG</u>	<u>A090718 001</u>	Mar 17, 2010
<u>AB</u>		<u>12.5MG; 15MG</u>	<u>A090718 002</u>	Mar 17, 2010
<u>AB</u>		<u>25MG; 15MG</u>	<u>A090718 003</u>	Mar 17, 2010
<u>AB</u>	TEVA	<u>12.5MG; 7.5MG</u>	<u>A076980 001</u>	Mar 07, 2007
<u>AB</u>		<u>12.5MG; 15MG</u>	<u>A076980 003</u>	Mar 07, 2007
<u>AB</u>	!	<u>25MG; 15MG</u>	<u>A076980 002</u>	Mar 07, 2007

HYDROCHLOROTHIAZIDE; OLMESARTAN MEDOXOMIL

TABLET; ORAL

BENICAR HCT

<u>AB</u>	+ COSETTE	<u>12.5MG; 20MG</u>	<u>N021532 002</u>	Jun 05, 2003
<u>AB</u>	+	<u>12.5MG; 40MG</u>	<u>N021532 003</u>	Jun 05, 2003
<u>AB</u>	+!	<u>25MG; 40MG</u>	<u>N021532 005</u>	Jun 05, 2003

HYDROCHLOROTHIAZIDE AND OLMESARTAN MEDOXOMIL

<u>AB</u>	ALKEM LABS LTD	<u>12.5MG; 20MG</u>	<u>A207037 001</u>	Jul 11, 2024
<u>AB</u>		<u>12.5MG; 40MG</u>	<u>A207037 002</u>	Jul 11, 2024
<u>AB</u>		<u>25MG; 40MG</u>	<u>A207037 003</u>	Jul 11, 2024

OLMESARTAN MEDOXOMIL AND HYDROCHLOROTHIAZIDE

<u>AB</u>	ACCORD HLTHCARE	<u>12.5MG; 20MG</u>	<u>A209281 001</u>	Feb 07, 2019
<u>AB</u>		<u>12.5MG; 40MG</u>	<u>A209281 002</u>	Feb 07, 2019
<u>AB</u>		<u>25MG; 40MG</u>	<u>A209281 003</u>	Feb 07, 2019
<u>AB</u>	ALEMBIC	<u>12.5MG; 20MG</u>	<u>A204233 001</u>	Apr 24, 2017
<u>AB</u>		<u>12.5MG; 40MG</u>	<u>A204233 002</u>	Apr 24, 2017
<u>AB</u>		<u>25MG; 40MG</u>	<u>A204233 003</u>	Apr 24, 2017
<u>AB</u>	AUROBINDO PHARMA	<u>12.5MG; 20MG</u>	<u>A205391 001</u>	Apr 24, 2017
<u>AB</u>		<u>12.5MG; 40MG</u>	<u>A205391 002</u>	Apr 24, 2017
<u>AB</u>		<u>25MG; 40MG</u>	<u>A205391 003</u>	Apr 24, 2017
<u>AB</u>	MACLEODS PHARMS LTD	<u>12.5MG; 20MG</u>	<u>A204801 001</u>	Jun 01, 2023
<u>AB</u>		<u>12.5MG; 40MG</u>	<u>A204801 002</u>	Jun 01, 2023
<u>AB</u>		<u>25MG; 40MG</u>	<u>A204801 003</u>	Jun 01, 2023
<u>AB</u>	PRINSTON INC	<u>12.5MG; 20MG</u>	<u>A207804 001</u>	Apr 24, 2017
<u>AB</u>		<u>12.5MG; 40MG</u>	<u>A207804 002</u>	Apr 24, 2017
<u>AB</u>		<u>25MG; 40MG</u>	<u>A207804 003</u>	Apr 24, 2017
<u>AB</u>	UMEDICA	<u>12.5MG; 20MG</u>	<u>A208847 001</u>	Sep 17, 2019
<u>AB</u>		<u>12.5MG; 40MG</u>	<u>A208847 003</u>	Sep 17, 2019
<u>AB</u>		<u>25MG; 40MG</u>	<u>A208847 002</u>	Sep 17, 2019

PRESCRIPTION DRUG PRODUCT LIST

HYDROCHLOROTHIAZIDE; QUINAPRIL HYDROCHLORIDE

TABLET; ORAL

QUINAPRIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE

AB	APOTEX	<u>12.5MG;EQ 10MG BASE</u>	<u>A091524 001</u>	Mar 12, 2013
AB		<u>12.5MG;EQ 20MG BASE</u>	<u>A091524 002</u>	Mar 12, 2013
AB		<u>25MG;EQ 20MG BASE</u>	<u>A091524 003</u>	Mar 12, 2013
AB	AUROBINDO PHARMA	<u>12.5MG;EQ 10MG BASE</u>	<u>A078450 001</u>	Aug 24, 2007
AB		<u>12.5MG;EQ 20MG BASE</u>	<u>A078450 002</u>	Aug 24, 2007
AB	!	<u>25MG;EQ 20MG BASE</u>	<u>A078450 003</u>	Aug 24, 2007
AB	CHARTWELL RX	<u>12.5MG;EQ 10MG BASE</u>	<u>A076374 001</u>	Mar 31, 2004
AB		<u>12.5MG;EQ 20MG BASE</u>	<u>A076374 002</u>	Mar 31, 2004
AB		<u>25MG;EQ 20MG BASE</u>	<u>A076374 003</u>	Mar 31, 2004

HYDROCHLOROTHIAZIDE; SPIRONOLACTONE

TABLET; ORAL

ALDACTAZIDE

AB	+ PFIZER	<u>25MG;25MG</u>	<u>N012616 004</u>	Dec 30, 1982
-----------	----------	------------------	--------------------	--------------

SPIRONOLACTONE AND HYDROCHLOROTHIAZIDE

AB	MYLAN	<u>25MG;25MG</u>	<u>A086513 001</u>	
AB	SUN PHARM INDUSTRIES	<u>25MG;25MG</u>	<u>A089534 001</u>	Jul 02, 1987

HYDROCHLOROTHIAZIDE; TELMISARTAN

TABLET; ORAL

MICARDIS HCT

AB	+ BOEHRINGER INGELHEIM	<u>12.5MG;40MG</u>	<u>N021162 001</u>	Nov 17, 2000
AB	+	<u>12.5MG;80MG</u>	<u>N021162 002</u>	Nov 17, 2000
AB	+	<u>25MG;80MG</u>	<u>N021162 003</u>	Apr 19, 2004

TELMISARTAN AND HYDROCHLOROTHIAZIDE

AB	ALEMBIC	<u>12.5MG;40MG</u>	<u>A203010 001</u>	Feb 25, 2014
AB		<u>12.5MG;80MG</u>	<u>A203010 002</u>	Feb 25, 2014
AB		<u>25MG;80MG</u>	<u>A203010 003</u>	Feb 25, 2014
AB	AUROBINDO PHARMA	<u>12.5MG;40MG</u>	<u>A208727 001</u>	Dec 15, 2016
AB		<u>12.5MG;80MG</u>	<u>A208727 002</u>	Dec 15, 2016
AB		<u>25MG;80MG</u>	<u>A208727 003</u>	Dec 15, 2016
AB	GLENMARK PHARMS LTD	<u>12.5MG;40MG</u>	<u>A202544 001</u>	Mar 04, 2019
AB		<u>12.5MG;80MG</u>	<u>A202544 002</u>	Mar 04, 2019
AB		<u>25MG;80MG</u>	<u>A202544 003</u>	Mar 04, 2019
AB	LUPIN LTD	<u>12.5MG;40MG</u>	<u>A091351 001</u>	Aug 07, 2014
AB		<u>12.5MG;80MG</u>	<u>A091351 002</u>	Aug 07, 2014
AB		<u>25MG;80MG</u>	<u>A091351 003</u>	Aug 07, 2014
AB	PRINSTON INC	<u>12.5MG;40MG</u>	<u>A209028 001</u>	Nov 06, 2017
AB		<u>12.5MG;80MG</u>	<u>A209028 002</u>	Nov 06, 2017
AB		<u>25MG;80MG</u>	<u>A209028 003</u>	Nov 06, 2017
AB	ZYDUS PHARMS	<u>12.5MG;40MG</u>	<u>A204221 001</u>	Aug 15, 2017
AB		<u>12.5MG;80MG</u>	<u>A204221 002</u>	Aug 15, 2017
AB		<u>25MG;80MG</u>	<u>A204221 003</u>	Aug 15, 2017

HYDROCHLOROTHIAZIDE; TRIAMTERENE

CAPSULE; ORAL

TRIAMTERENE AND HYDROCHLOROTHIAZIDE

AB	CADILA	<u>25MG;37.5MG</u>	<u>A208358 001</u>	Feb 11, 2019
AB	LANNETT CO INC	<u>25MG;37.5MG</u>	<u>A201407 001</u>	Dec 09, 2011
AB	! SANDOZ	<u>25MG;37.5MG</u>	<u>A074821 001</u>	Jun 05, 1997

TABLET; ORAL

MAXZIDE

AB	+! AUROBINDO PHARMA USA	<u>50MG;75MG</u>	<u>N019129 001</u>	Oct 22, 1984
-----------	-------------------------	------------------	--------------------	--------------

MAXZIDE-25

AB	+ AUROBINDO PHARMA USA	<u>25MG;37.5MG</u>	<u>N019129 003</u>	May 13, 1988
-----------	------------------------	--------------------	--------------------	--------------

TRIAMTERENE AND HYDROCHLOROTHIAZIDE

AB	APOTEX INC	<u>25MG;37.5MG</u>	<u>A071251 002</u>	May 05, 1998
AB		<u>50MG;75MG</u>	<u>A071251 001</u>	Apr 17, 1988
AB	CHARTWELL RX	<u>50MG;75MG</u>	<u>A072011 001</u>	Jun 17, 1988
AB	RUBICON	<u>25MG;37.5MG</u>	<u>A216211 001</u>	Feb 23, 2022
AB		<u>50MG;75MG</u>	<u>A216211 002</u>	Feb 23, 2022
AB	SANDOZ	<u>25MG;37.5MG</u>	<u>A073281 001</u>	Apr 30, 1992
AB	WATSON LABS	<u>25MG;37.5MG</u>	<u>A073449 001</u>	Sep 23, 1993
AB		<u>50MG;75MG</u>	<u>A071851 001</u>	Nov 30, 1988
AB	ZYDUS PHARMS	<u>25MG;37.5MG</u>	<u>A208360 001</u>	Jun 29, 2018
AB		<u>50MG;75MG</u>	<u>A208360 002</u>	Jun 29, 2018

PRESCRIPTION DRUG PRODUCT LIST

HYDROCHLOROTHIAZIDE; VALSARTAN

TABLET; ORAL

DIOVAN HCT

<u>AB</u>	+	NOVARTIS	<u>12.5MG;80MG</u>	<u>N020818</u>	<u>001</u>	Mar 06, 1998
<u>AB</u>	+		<u>12.5MG;160MG</u>	<u>N020818</u>	<u>002</u>	Mar 06, 1998
<u>AB</u>	+		<u>12.5MG;320MG</u>	<u>N020818</u>	<u>004</u>	Apr 28, 2006
<u>AB</u>	+		<u>25MG;160MG</u>	<u>N020818</u>	<u>003</u>	Jan 17, 2002
<u>AB</u>	+	!	<u>25MG;320MG</u>	<u>N020818</u>	<u>005</u>	Apr 28, 2006

VALSARTAN AND HYDROCHLOROTHIAZIDE

<u>AB</u>		ALEMBIC	<u>12.5MG;80MG</u>	<u>A201662</u>	<u>001</u>	Mar 21, 2013
<u>AB</u>			<u>12.5MG;160MG</u>	<u>A201662</u>	<u>002</u>	Mar 21, 2013
<u>AB</u>			<u>12.5MG;320MG</u>	<u>A201662</u>	<u>003</u>	Mar 21, 2013
<u>AB</u>			<u>25MG;160MG</u>	<u>A201662</u>	<u>004</u>	Mar 21, 2013
<u>AB</u>			<u>25MG;320MG</u>	<u>A201662</u>	<u>005</u>	Mar 21, 2013
<u>AB</u>		AMNEAL PHARMS	<u>12.5MG;80MG</u>	<u>A204382</u>	<u>001</u>	Aug 11, 2023
<u>AB</u>			<u>12.5MG;160MG</u>	<u>A204382</u>	<u>002</u>	Aug 11, 2023
<u>AB</u>			<u>12.5MG;320MG</u>	<u>A204382</u>	<u>004</u>	Aug 11, 2023
<u>AB</u>			<u>25MG;160MG</u>	<u>A204382</u>	<u>003</u>	Aug 11, 2023
<u>AB</u>			<u>25MG;320MG</u>	<u>A204382</u>	<u>005</u>	Aug 11, 2023
<u>AB</u>		AUROBINDO PHARMA LTD	<u>12.5MG;80MG</u>	<u>A202519</u>	<u>001</u>	Mar 21, 2013
<u>AB</u>			<u>12.5MG;160MG</u>	<u>A202519</u>	<u>002</u>	Mar 21, 2013
<u>AB</u>			<u>12.5MG;320MG</u>	<u>A202519</u>	<u>003</u>	Mar 21, 2013
<u>AB</u>			<u>25MG;160MG</u>	<u>A202519</u>	<u>004</u>	Mar 21, 2013
<u>AB</u>			<u>25MG;320MG</u>	<u>A202519</u>	<u>005</u>	Mar 21, 2013
<u>AB</u>		LUPIN LTD	<u>12.5MG;80MG</u>	<u>A078946</u>	<u>003</u>	Mar 21, 2013
<u>AB</u>			<u>12.5MG;160MG</u>	<u>A078946</u>	<u>004</u>	Mar 21, 2013
<u>AB</u>			<u>12.5MG;320MG</u>	<u>A078946</u>	<u>001</u>	Mar 21, 2013
<u>AB</u>			<u>25MG;160MG</u>	<u>A078946</u>	<u>005</u>	Mar 21, 2013
<u>AB</u>			<u>25MG;320MG</u>	<u>A078946</u>	<u>002</u>	Mar 21, 2013
<u>AB</u>		MACLEODS PHARMS LTD	<u>12.5MG;80MG</u>	<u>A203145</u>	<u>001</u>	Apr 19, 2013
<u>AB</u>			<u>12.5MG;160MG</u>	<u>A203145</u>	<u>002</u>	Apr 19, 2013
<u>AB</u>			<u>12.5MG;320MG</u>	<u>A203145</u>	<u>003</u>	Apr 19, 2013
<u>AB</u>			<u>25MG;160MG</u>	<u>A203145</u>	<u>004</u>	Apr 19, 2013
<u>AB</u>			<u>25MG;320MG</u>	<u>A203145</u>	<u>005</u>	Apr 19, 2013
<u>AB</u>		MYLAN PHARMS INC	<u>12.5MG;80MG</u>	<u>A078020</u>	<u>001</u>	Sep 21, 2012
<u>AB</u>			<u>12.5MG;160MG</u>	<u>A078020</u>	<u>002</u>	Sep 21, 2012
<u>AB</u>			<u>12.5MG;320MG</u>	<u>A078020</u>	<u>004</u>	Sep 21, 2012
<u>AB</u>			<u>25MG;160MG</u>	<u>A078020</u>	<u>003</u>	Sep 21, 2012
<u>AB</u>			<u>25MG;320MG</u>	<u>A078020</u>	<u>005</u>	Sep 21, 2012
<u>AB</u>		PRINSTON INC	<u>12.5MG;80MG</u>	<u>A206083</u>	<u>001</u>	Feb 08, 2016
<u>AB</u>			<u>12.5MG;160MG</u>	<u>A206083</u>	<u>002</u>	Feb 08, 2016
<u>AB</u>			<u>12.5MG;320MG</u>	<u>A206083</u>	<u>003</u>	Feb 08, 2016
<u>AB</u>			<u>25MG;160MG</u>	<u>A206083</u>	<u>004</u>	Feb 08, 2016
<u>AB</u>			<u>25MG;320MG</u>	<u>A206083</u>	<u>005</u>	Feb 08, 2016

HYDROCODONE BITARTRATE

CAPSULE, EXTENDED RELEASE; ORAL

HYDROCODONE BITARTRATE

!	ALVOGEN	10MG	A206986	001	Jan 21, 2020
		15MG	A206986	002	Jan 21, 2020
		20MG	A206986	003	Jan 21, 2020
		30MG	A206986	004	Jan 21, 2020
		40MG	A206986	005	Jan 21, 2020
		50MG	A206986	006	Jan 21, 2020

TABLET, EXTENDED RELEASE; ORAL

HYDROCODONE BITARTRATE

<u>AB</u>		ALVOGEN	<u>20MG</u>	<u>A208269</u>	<u>001</u>	Mar 01, 2021
<u>AB</u>			<u>30MG</u>	<u>A208269</u>	<u>002</u>	Mar 01, 2021
<u>AB</u>			<u>40MG</u>	<u>A208269</u>	<u>003</u>	Mar 01, 2021
<u>AB</u>			<u>60MG</u>	<u>A208269</u>	<u>004</u>	Mar 01, 2021
<u>AB</u>			<u>80MG</u>	<u>A208269</u>	<u>005</u>	Mar 01, 2021
<u>AB</u>			<u>100MG</u>	<u>A208269</u>	<u>006</u>	Mar 01, 2021
		<u>HYSINGLA ER</u>				
<u>AB</u>	+	! PURDUE PHARMA LP	<u>20MG</u>	<u>N206627</u>	<u>001</u>	Nov 20, 2014
<u>AB</u>	+		<u>30MG</u>	<u>N206627</u>	<u>002</u>	Nov 20, 2014
<u>AB</u>	+		<u>40MG</u>	<u>N206627</u>	<u>003</u>	Nov 20, 2014
<u>AB</u>	+		<u>60MG</u>	<u>N206627</u>	<u>004</u>	Nov 20, 2014
<u>AB</u>	+		<u>80MG</u>	<u>N206627</u>	<u>005</u>	Nov 20, 2014
<u>AB</u>	+		<u>100MG</u>	<u>N206627</u>	<u>006</u>	Nov 20, 2014
		HYDROCODONE BITARTRATE				
		ALVOGEN	120MG	A208269	007	Mar 01, 2021

PRESCRIPTION DRUG PRODUCT LIST

HYDROCODONE BITARTRATE; IBUPROFEN

TABLET; ORAL

HYDROCODONE BITARTRATE AND IBUPROFEN

<u>AB</u>	ACTAVIS LABS FL INC	<u>7.5MG;200MG</u>	<u>A076604 001</u>	Dec 31, 2003
<u>AB</u>	! AMNEAL PHARMS NY	<u>7.5MG;200MG</u>	<u>A076642 001</u>	Oct 12, 2004
<u>AB</u>	AUROLIFE PHARMA LLC	<u>7.5MG;200MG</u>	<u>A204575 001</u>	Jun 02, 2016
	AMNEAL PHARMS NY	5MG;200MG	A076642 002	Mar 18, 2004

HYDROCORTISONE

CREAM; TOPICAL

ALA-CORT

<u>AT</u>	CROWN LABS	<u>2.5%</u>	<u>A080706 007</u>	Jan 05, 2016
<u>AT</u>		<u>1%</u>	<u>A080706 006</u>	

ANUSOL HC

<u>AT</u>	SALIX PHARMS	<u>2.5%</u>	<u>A088250 001</u>	Jun 06, 1984
-----------	--------------	-------------	--------------------	--------------

HYDROCORTISONE

<u>AT</u>	ACTAVIS MID ATLANTIC	<u>1%</u>	<u>A087795 001</u>	May 03, 1983
<u>AT</u>		<u>2.5%</u>	<u>A089682 001</u>	Mar 10, 1988
<u>AT</u>	+! FOUGERA PHARMS INC	<u>1%</u>	<u>A080693 003</u>	
<u>AT</u>	!	<u>2.5%</u>	<u>A089414 001</u>	Dec 16, 1986
<u>AT</u>	PADAGIS US	<u>2.5%</u>	<u>A085025 001</u>	
<u>AT</u>	RISING	<u>2.5%</u>	<u>A040879 001</u>	Aug 20, 2010
<u>AT</u>	TARO	<u>2.5%</u>	<u>A088799 001</u>	Nov 09, 1984

ENEMA; RECTAL

COLOCORT

<u>AB</u>	CHARTWELL	<u>100MG/60ML</u>	<u>A075172 001</u>	Dec 03, 1999
-----------	-----------	-------------------	--------------------	--------------

CORTENEMA

<u>AB</u>	+! ANI PHARMS	<u>100MG/60ML</u>	<u>N016199 001</u>	
-----------	---------------	-------------------	--------------------	--

GRANULE; ORAL

ALKINDI SPRINKLE

+	ETON	0.5MG	N213876 001	Sep 29, 2020
+		1MG	N213876 002	Sep 29, 2020
+		2MG	N213876 003	Sep 29, 2020
+	!	5MG	N213876 004	Sep 29, 2020

LOTION; TOPICAL

HYDROCORTISONE

<u>AT</u>	TARO	<u>2.5%</u>	<u>A040247 001</u>	Jul 23, 1999
-----------	------	-------------	--------------------	--------------

STIE-CORT

<u>AT</u>	! PADAGIS US	<u>2.5%</u>	<u>A089074 001</u>	Nov 26, 1985
-----------	--------------	-------------	--------------------	--------------

ALA-SCALP

	LEGACY PHARMA	2%	A083231 001	
--	---------------	----	-------------	--

OINTMENT; TOPICAL

HYDROCORTISONE

<u>AT</u>	+! FOUGERA PHARMS	<u>1%</u>	<u>A080692 001</u>	
<u>AT</u>	! FOUGERA PHARMS INC	<u>2.5%</u>	<u>A081203 001</u>	May 28, 1993
<u>AT</u>	PADAGIS US	<u>2.5%</u>	<u>A085027 001</u>	
<u>AT</u>	TARO	<u>1%</u>	<u>A086257 001</u>	

HYDROCORTISONE IN ABSORBASE

<u>AT</u>	CMP PHARMA INC	<u>1%</u>	<u>A088138 001</u>	Sep 06, 1985
-----------	----------------	-----------	--------------------	--------------

SOLUTION; TOPICAL

TEXACORT

!	MISSION PHARMA	2.5%	A081271 001	Apr 17, 1992
---	----------------	------	-------------	--------------

TABLET; ORAL

CORTEF

<u>AB</u>	+ PHARMACIA AND UPJOHN	<u>5MG</u>	<u>N008697 003</u>	
-----------	------------------------	------------	--------------------	--

<u>AB</u>	+	<u>10MG</u>	<u>N008697 001</u>	
-----------	---	-------------	--------------------	--

<u>AB</u>	+!	<u>20MG</u>	<u>N008697 002</u>	
-----------	----	-------------	--------------------	--

HYDROCORTISONE

<u>AB</u>	AUROBINDO PHARMA LTD	<u>5MG</u>	<u>A214649 001</u>	Jul 17, 2023
<u>AB</u>		<u>10MG</u>	<u>A214649 002</u>	Jul 17, 2023
<u>AB</u>		<u>20MG</u>	<u>A214649 003</u>	Jul 17, 2023
<u>AB</u>	HIBROW HLTHCARE	<u>5MG</u>	<u>A217160 001</u>	Nov 25, 2024
<u>AB</u>		<u>10MG</u>	<u>A217160 002</u>	Nov 25, 2024
<u>AB</u>		<u>20MG</u>	<u>A217160 003</u>	Nov 25, 2024
<u>AB</u>	IMPAX LABS INC	<u>5MG</u>	<u>A040646 001</u>	Mar 30, 2007
<u>AB</u>		<u>10MG</u>	<u>A040646 002</u>	Mar 30, 2007
<u>AB</u>		<u>20MG</u>	<u>A040646 003</u>	Mar 30, 2007
<u>AB</u>	STRIDES PHARMA	<u>5MG</u>	<u>A207029 001</u>	Apr 27, 2017
<u>AB</u>		<u>10MG</u>	<u>A207029 002</u>	Apr 27, 2017
<u>AB</u>		<u>20MG</u>	<u>A207029 003</u>	Apr 27, 2017

PRESCRIPTION DRUG PRODUCT LIST

HYDROCORTISONE ACETATE

AEROSOL, METERED; RECTAL

CORTIFOAM

+! MYLAN SPECIALITY LP 10%

N017351 001 Feb 10, 1982

CREAM; TOPICAL

MICORT-HC

LEGACY PHARMA 2.5%

A040396 001 Feb 27, 2001

HYDROCORTISONE ACETATE; PRAMOXINE HYDROCHLORIDE

AEROSOL, METERED; TOPICAL

EPIFOAM

BX MYLAN SPECIALITY LP 1%;1%

A086457 001

PROCTOFOAM HC

BX +! MYLAN SPECIALITY LP 1%;1%

A086195 001

CREAM; TOPICAL

PRAMOSONE

LEGACY PHARMA 0.5%;1%

A083778 001

1%;1%

A085368 001

LOTION; TOPICAL

PRAMOSONE

LEGACY PHARMA 1%;1%

A085980 001

2.5%;1%

A085979 001

HYDROCORTISONE BUTYRATE

CREAM; TOPICAL

HYDROCORTISONE BUTYRATE**AB1** TARO PHARM INDS **0.1%****A076654 001** Aug 03, 2005LOCOID**AB1** +! BAUSCH **0.1%****N018514 001** Mar 31, 1982HYDROCORTISONE BUTYRATE**AB2** ACTAVIS MID **0.1%****A205134 001** Dec 08, 2017

ATLANTIC

AB2 GLENMARK PHARMS LTD **0.1%****A202145 001** Sep 27, 2013LOCOID LIPOCREAM**AB2** +! PRECISION DERMAT **0.1%****N020769 001** Sep 08, 1997

LOTION; TOPICAL

HYDROCORTISONE BUTYRATE**AB** ! LUPIN LTD **0.1%****A210209 001** Aug 17, 2018**AB** THE J MOLNER **0.1%****A209556 001** Nov 21, 2017

OINTMENT; TOPICAL

HYDROCORTISONE BUTYRATE**AB** TARO **0.1%****A076842 001** Dec 27, 2004LOCOID**AB** +! PRECISION DERMAT **0.1%****N018652 001** Oct 29, 1982

SOLUTION; TOPICAL

HYDROCORTISONE BUTYRATE**AT** TARO PHARM INDS **0.1%****A076364 001** Jan 14, 2004LOCOID**AT** +! BAUSCH **0.1%****N019116 001** Feb 25, 1987HYDROCORTISONE SODIUM SUCCINATE

INJECTABLE; INJECTION

HYDROCORTISONE SODIUM SUCCINATE**AP** CIPLA **EQ 100MG BASE/VIAL****A214050 001** Sep 05, 2024SOLU-CORTEF**AP** +! PHARMACIA AND **EQ 100MG BASE/VIAL****N009866 001**

UPJOHN

+! EQ 250MG BASE/VIAL

N009866 002

+! EQ 500MG BASE/VIAL

N009866 003

+! EQ 1GM BASE/VIAL

N009866 004

HYDROCORTISONE VALERATE

CREAM; TOPICAL

HYDROCORTISONE VALERATE**AB** COSETTE **0.2%****A213724 001** Feb 11, 2021**AB** ENCUBE ETHICALS **0.2%****A211047 001** Nov 04, 2021**AB** GLENMARK PHARMS LTD **0.2%****A211129 001** Oct 12, 2018**AB** LUPIN LTD **0.2%****A210307 001** Aug 15, 2019**AB** PADAGIS ISRAEL **0.2%****A075666 001** May 24, 2000**AB** ! TARO **0.2%****A075042 001** Aug 25, 1998

OINTMENT; TOPICAL

HYDROCORTISONE VALERATE**AB** COSETTE **0.2%****A211764 001** Mar 04, 2020**AB** GLENMARK PHARMS LTD **0.2%****A211750 001** Dec 14, 2018**AB** ! TARO **0.2%****A075043 001** Aug 25, 1998

PRESCRIPTION DRUG PRODUCT LIST

HYDROCORTISONE; NEOMYCIN SULFATE; POLYMYXIN B SULFATE

SOLUTION/DROPS;OTIC

NEOMYCIN AND POLYMYXIN B SULFATES AND HYDROCORTISONE

<u>AT</u>	!	BAUSCH AND LOMB	<u>1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML</u>	<u>A064053 001</u>	Dec 29, 1995
<u>AT</u>		SANDOZ	<u>1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML</u>	<u>A062423 001</u>	Aug 25, 1983

SUSPENSION/DROPS;OPHTHALMIC

NEOMYCIN AND POLYMYXIN B SULFATES AND HYDROCORTISONE

!	SANDOZ	1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML	A062874 001	May 11, 1988
---	--------	-------------------------------------	-------------	--------------

SUSPENSION/DROPS;OTIC

CASPORYN HC

<u>AT</u>	+	CASPER PHARMA LLC	<u>1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML</u>	<u>N060613 001</u>	
-----------	---	-------------------	--	--------------------	--

NEOMYCIN AND POLYMYXIN B SULFATES AND HYDROCORTISONE

<u>AT</u>		AMRING PHARMS	<u>1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML</u>	<u>A065219 001</u>	May 01, 2006
<u>AT</u>		SANDOZ	<u>1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML</u>	<u>A062488 001</u>	Nov 06, 1985

OTICAIR

<u>AT</u>		BAUSCH AND LOMB	<u>1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML</u>	<u>A064065 001</u>	Aug 28, 1996
-----------	--	-----------------	--	--------------------	--------------

HYDROMORPHONE HYDROCHLORIDE

INJECTABLE;INJECTION

DILAUDID

<u>AP</u>	+	FRESENIUS KABI USA	<u>0.2MG/ML</u>	<u>N019034 006</u>	Jan 16, 2020
<u>AP</u>	+		<u>0.5MG/0.5ML</u>	<u>N019034 007</u>	Feb 10, 2017
<u>AP</u>	+		<u>1MG/ML</u>	<u>N019034 003</u>	Apr 30, 2009
<u>AP</u>	+		<u>2MG/ML</u>	<u>N019034 004</u>	Apr 30, 2009
<u>AP</u>	+		<u>4MG/ML</u>	<u>N019034 005</u>	Apr 30, 2009

HYDROMORPHONE HYDROCHLORIDE

<u>AP</u>		HIKMA	<u>0.2MG/ML</u>	<u>A216899 001</u>	Feb 09, 2024
<u>AP</u>			<u>0.5MG/0.5ML</u>	<u>A216899 002</u>	Feb 09, 2024
<u>AP</u>			<u>1MG/ML</u>	<u>A216899 003</u>	Feb 09, 2024
<u>AP</u>			<u>2MG/ML</u>	<u>A202159 001</u>	Apr 27, 2018
<u>AP</u>			<u>2MG/ML</u>	<u>A216899 004</u>	Feb 09, 2024
<u>AP</u>		HOSPIRA INC	<u>1MG/ML</u>	<u>N200403 001</u>	Dec 01, 2011
<u>AP</u>			<u>2MG/ML</u>	<u>N200403 002</u>	Dec 01, 2011
<u>AP</u>			<u>4MG/ML</u>	<u>N200403 003</u>	Dec 01, 2011
<u>AP</u>			<u>10MG/ML</u>	<u>A078591 001</u>	Jun 17, 2008
<u>AP</u>		RISING	<u>10MG/ML</u>	<u>A078228 001</u>	Apr 14, 2010
<u>AP</u>			<u>10MG/ML</u>	<u>A078261 001</u>	Apr 14, 2010
		HOSPIRA INC	0.25MG/0.5ML	N200403 005	Mar 03, 2023
			0.5MG/0.5ML	N200403 004	Jun 13, 2022

SOLUTION;INTRAMUSCULAR, INTRAVENOUS, SUBCUTANEOUS

HYDROMORPHONE HYDROCHLORIDE

+	HIKMA	40MG/20ML (2MG/ML)	N217812 001	Dec 14, 2023
---	-------	--------------------	-------------	--------------

SOLUTION;ORAL

DILAUDID

<u>AA</u>	+	RHODES PHARMS	<u>5MG/5ML</u>	<u>N019891 001</u>	Dec 07, 1992
-----------	---	---------------	----------------	--------------------	--------------

HYDROMORPHONE HYDROCHLORIDE

<u>AA</u>		ASCENT PHARMS INC	<u>5MG/5ML</u>	<u>A210176 001</u>	Oct 27, 2017
<u>AA</u>		HIKMA	<u>5MG/5ML</u>	<u>A074653 001</u>	Jul 29, 1998

TABLET;ORAL

DILAUDID

<u>AB</u>	+	RHODES PHARMS	<u>2MG</u>	<u>N019892 003</u>	Nov 09, 2007
<u>AB</u>	+		<u>4MG</u>	<u>N019892 002</u>	Nov 09, 2007
<u>AB</u>	+		<u>8MG</u>	<u>N019892 001</u>	Dec 07, 1992

HYDROMORPHONE HYDROCHLORIDE

<u>AB</u>		ASCENT PHARMS INC	<u>2MG</u>	<u>A210506 001</u>	Jan 17, 2018
<u>AB</u>			<u>4MG</u>	<u>A210506 002</u>	Jan 17, 2018
<u>AB</u>			<u>8MG</u>	<u>A210506 003</u>	Jan 17, 2018
<u>AB</u>		AUROLIFE PHARMA LLC	<u>2MG</u>	<u>A205814 001</u>	May 13, 2016
<u>AB</u>			<u>4MG</u>	<u>A205814 002</u>	May 13, 2016
<u>AB</u>			<u>8MG</u>	<u>A205814 003</u>	May 13, 2016
<u>AB</u>		SPECGX LLC	<u>2MG</u>	<u>A076855 002</u>	Sep 19, 2007
<u>AB</u>			<u>4MG</u>	<u>A076855 003</u>	Sep 19, 2007
<u>AB</u>			<u>8MG</u>	<u>A076855 001</u>	Dec 23, 2004

TABLET, EXTENDED RELEASE;ORAL

HYDROMORPHONE HYDROCHLORIDE

<u>AB</u>		ASCENT PHARMS INC	<u>8MG</u>	<u>A212133 001</u>	Sep 23, 2020
<u>AB</u>			<u>12MG</u>	<u>A212133 002</u>	Sep 23, 2020
<u>AB</u>			<u>16MG</u>	<u>A212133 003</u>	Sep 23, 2020
<u>AB</u>			<u>32MG</u>	<u>A212133 004</u>	Sep 23, 2020
<u>AB</u>		OSMOTICA PHARM US	<u>8MG</u>	<u>A205629 001</u>	Jul 07, 2016
<u>AB</u>			<u>12MG</u>	<u>A205629 002</u>	Jul 07, 2016
<u>AB</u>			<u>16MG</u>	<u>A205629 003</u>	Jul 07, 2016
<u>AB</u>	!		<u>32MG</u>	<u>A205629 004</u>	Jul 07, 2016

PRESCRIPTION DRUG PRODUCT LIST

HYDROMORPHONE HYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL

HYDROMORPHONE HYDROCHLORIDE

<u>AB</u>	PADAGIS US	<u>8MG</u>	<u>A204278</u>	<u>001</u>	Apr 06, 2015
<u>AB</u>		<u>12MG</u>	<u>A204278</u>	<u>002</u>	Apr 06, 2015
<u>AB</u>		<u>16MG</u>	<u>A204278</u>	<u>003</u>	Apr 06, 2015
<u>AB</u>		<u>32MG</u>	<u>A204278</u>	<u>004</u>	Sep 20, 2017
	OSMOTICA PHARM US	24MG	A205629	005	Oct 30, 2024

HYDROXOCOBALAMIN

INJECTABLE; INJECTION

CYANOKIT

+! BTG INTL 5GM/VIAL (5GM/KIT) N022041 001 Apr 08, 2011

HYDROXOCOBALAMIN

! ACTAVIS 1MG/ML A085998 001

HYDROXYCHLOROQUINE SULFATE

TABLET; ORAL

HYDROXYCHLOROQUINE SULFATE

<u>AB</u>	ACCORD HLTHCARE	<u>100MG</u>	<u>A213342</u>	<u>002</u>	Aug 18, 2021
<u>AB</u>		<u>200MG</u>	<u>A213342</u>	<u>001</u>	Apr 07, 2020
<u>AB</u>		<u>300MG</u>	<u>A213342</u>	<u>003</u>	Aug 18, 2021
<u>AB</u>	!	<u>400MG</u>	<u>A213342</u>	<u>004</u>	Aug 18, 2021
<u>AB</u>	ALKALOIDA ZRT	<u>200MG</u>	<u>A201691</u>	<u>001</u>	May 08, 2018
<u>AB</u>	AMNEAL PHARMS CO	<u>200MG</u>	<u>A210577</u>	<u>001</u>	May 15, 2018
<u>AB</u>	!	<u>100MG</u>	<u>A210441</u>	<u>002</u>	Sep 19, 2022
<u>AB</u>		<u>200MG</u>	<u>A210441</u>	<u>001</u>	May 01, 2018
<u>AB</u>		<u>300MG</u>	<u>A210441</u>	<u>003</u>	Sep 19, 2022
<u>AB</u>		<u>400MG</u>	<u>A210441</u>	<u>004</u>	Sep 19, 2022
<u>AB</u>	AUROBINDO PHARMA USA	<u>200MG</u>	<u>A040274</u>	<u>001</u>	May 29, 1998
<u>AB</u>	CHARTWELL RX	<u>200MG</u>	<u>A210543</u>	<u>001</u>	Jul 06, 2018
<u>AB</u>	CREEKWOOD PHARMS	<u>200MG</u>	<u>A040150</u>	<u>001</u>	Jan 27, 1996
<u>AB</u>	IPCA LABS LTD	<u>200MG</u>	<u>A040766</u>	<u>001</u>	Jun 14, 2007
<u>AB</u>	LAURUS	<u>200MG</u>	<u>A210959</u>	<u>001</u>	Jan 15, 2019
<u>AB</u>	SANDOZ	<u>200MG</u>	<u>A040104</u>	<u>001</u>	Nov 30, 1995
<u>AB</u>	SENORES PHARMS	<u>200MG</u>	<u>A212902</u>	<u>001</u>	May 14, 2020
<u>AB</u>	TEVA PHARMS	<u>200MG</u>	<u>A040081</u>	<u>001</u>	Sep 30, 1994
<u>AB</u>	ZYDUS PHARMS USA INC	<u>200MG</u>	<u>A040657</u>	<u>001</u>	Sep 21, 2007

PLAQUENIL

<u>AB</u>	+! ADVANZ PHARMA	<u>200MG</u>	<u>N009768</u>	<u>001</u>	
	SOVUNA				
	+ NOVITIUM PHARMA	200MG	N214581	001	Jan 14, 2022
	+!	300MG	N214581	002	Jan 14, 2022

HYDROXYUREA

CAPSULE; ORAL

HYDREA

<u>AB</u>	+! CHEPLAPHARM	<u>500MG</u>	<u>N016295</u>	<u>001</u>	
	<u>HYDROXYUREA</u>				
<u>AB</u>	BARR	<u>500MG</u>	<u>A075143</u>	<u>001</u>	Oct 16, 1998
<u>AB</u>	ENDO OPERATIONS	<u>500MG</u>	<u>A075340</u>	<u>001</u>	Feb 24, 1999
<u>AB</u>	LEADING	<u>500MG</u>	<u>A213438</u>	<u>001</u>	Apr 08, 2020
<u>AB</u>	QILU	<u>500MG</u>	<u>A218021</u>	<u>001</u>	Mar 06, 2024

SOLUTION; ORAL

XROMI

+! NOVA LABS LTD 100MG/ML N216593 001 Apr 04, 2024

TABLET; ORAL

SIKLOS

+ THERAVIA 100MG N208843 001 Dec 21, 2017

+! 1GM N208843 002 Dec 21, 2017

HYDROXYZINE HYDROCHLORIDE

INJECTABLE; INJECTION

HYDROXYZINE HYDROCHLORIDE

+! AM REGENT 25MG/ML A087408 001

+! 50MG/ML A087408 002

SYRUP; ORAL

HYDROXYZINE HYDROCHLORIDE

<u>AA</u>	APOZEAL PHARMS	<u>10MG/5ML</u>	<u>A210634</u>	<u>001</u>	Feb 26, 2019
<u>AA</u>	+! CHARTWELL RX	<u>10MG/5ML</u>	<u>A087294</u>	<u>001</u>	Apr 12, 1982
<u>AA</u>	LANNETT CO INC	<u>10MG/5ML</u>	<u>A201674</u>	<u>001</u>	Aug 21, 2013
<u>AA</u>	PAI HOLDINGS PHARM	<u>10MG/5ML</u>	<u>A040391</u>	<u>001</u>	Apr 10, 2002

PRESCRIPTION DRUG PRODUCT LIST

HYDROXYZINE HYDROCHLORIDE

TABLET; ORAL

HYDROXYZINE HYDROCHLORIDE

<u>AB</u>	AMNEAL PHARM	<u>10MG</u>	<u>A040808</u>	<u>001</u>	Sep 24, 2008
<u>AB</u>		<u>25MG</u>	<u>A040808</u>	<u>002</u>	Sep 24, 2008
<u>AB</u>		<u>50MG</u>	<u>A040808</u>	<u>003</u>	Sep 24, 2008
<u>AB</u>	AUROBINDO PHARMA LTD	<u>10MG</u>	<u>A087871</u>	<u>002</u>	Dec 20, 1982
<u>AB</u>		<u>25MG</u>	<u>A087871</u>	<u>003</u>	Dec 20, 1982
<u>AB</u>		<u>50MG</u>	<u>A087871</u>	<u>001</u>	Dec 20, 1982
<u>AB</u>	CHARTWELL RX	<u>10MG</u>	<u>A040804</u>	<u>001</u>	Jun 30, 2008
<u>AB</u>		<u>25MG</u>	<u>A040804</u>	<u>002</u>	Jun 30, 2008
<u>AB</u>		<u>50MG</u>	<u>A040804</u>	<u>003</u>	Jun 30, 2008
<u>AB</u>	EPIC PHARMA LLC	<u>10MG</u>	<u>A040604</u>	<u>002</u>	Dec 28, 2004
<u>AB</u>		<u>25MG</u>	<u>A040604</u>	<u>003</u>	Dec 28, 2004
<u>AB</u>		<u>50MG</u>	<u>A040604</u>	<u>001</u>	Dec 28, 2004
<u>AB</u>	GRAVITI PHARMS	<u>10MG</u>	<u>A217652</u>	<u>001</u>	Aug 17, 2023
<u>AB</u>		<u>25MG</u>	<u>A217652</u>	<u>002</u>	Aug 17, 2023
<u>AB</u>		<u>50MG</u>	<u>A217652</u>	<u>003</u>	Aug 17, 2023
<u>AB</u>	HERITAGE PHARMA	<u>10MG</u>	<u>A204279</u>	<u>001</u>	Aug 20, 2014
<u>AB</u>		<u>25MG</u>	<u>A204279</u>	<u>002</u>	Aug 20, 2014
<u>AB</u>		<u>50MG</u>	<u>A204279</u>	<u>003</u>	Aug 20, 2014
<u>AB</u>	HETERO LABS LTD III	<u>10MG</u>	<u>A040805</u>	<u>001</u>	May 29, 2008
<u>AB</u>		<u>25MG</u>	<u>A040805</u>	<u>002</u>	May 29, 2008
<u>AB</u>		<u>50MG</u>	<u>A040805</u>	<u>003</u>	May 29, 2008
<u>AB</u>	KVK TECH	<u>10MG</u>	<u>A040786</u>	<u>001</u>	Mar 20, 2007
<u>AB</u>		<u>25MG</u>	<u>A040786</u>	<u>002</u>	Mar 20, 2007
<u>AB</u>		<u>50MG</u>	<u>A040786</u>	<u>003</u>	Mar 20, 2007
<u>AB</u>	NORTHSTAR HLTHCARE	<u>10MG</u>	<u>A040840</u>	<u>002</u>	Mar 31, 2008
<u>AB</u>		<u>25MG</u>	<u>A040840</u>	<u>003</u>	Mar 31, 2008
<u>AB</u>		<u>50MG</u>	<u>A040840</u>	<u>001</u>	Mar 31, 2008
<u>AB</u>	NUVO PHARMS INC	<u>10MG</u>	<u>A207121</u>	<u>002</u>	Mar 29, 2017
<u>AB</u>		<u>25MG</u>	<u>A207121</u>	<u>001</u>	Mar 29, 2017
<u>AB</u>		<u>50MG</u>	<u>A207121</u>	<u>003</u>	Mar 29, 2017
<u>AB</u>	+! PLIVA	<u>10MG</u>	<u>A088617</u>	<u>001</u>	Jan 10, 1986
<u>AB</u>	+!	<u>25MG</u>	<u>A088618</u>	<u>001</u>	Jan 10, 1986
<u>AB</u>	+!	<u>50MG</u>	<u>A088619</u>	<u>001</u>	Jan 10, 1986
<u>AB</u>	PRINSTON INC	<u>10MG</u>	<u>A040580</u>	<u>003</u>	May 27, 2005
<u>AB</u>		<u>25MG</u>	<u>A040580</u>	<u>002</u>	May 27, 2005
<u>AB</u>		<u>50MG</u>	<u>A040580</u>	<u>001</u>	May 27, 2005

HYDROXYZINE PAMOATE

CAPSULE; ORAL

HYDROXYZINE PAMOATE

<u>AB</u>	BARR	<u>EQ 25MG HYDROCHLORIDE</u>	<u>A088496</u>	<u>001</u>	Jun 15, 1984
<u>AB</u>		<u>EQ 50MG HYDROCHLORIDE</u>	<u>A088487</u>	<u>001</u>	Jun 15, 1984
<u>AB</u>	HERITAGE PHARMA	<u>EQ 25MG HYDROCHLORIDE</u>	<u>A201507</u>	<u>001</u>	Jun 03, 2013
<u>AB</u>	!	<u>EQ 50MG HYDROCHLORIDE</u>	<u>A201507</u>	<u>002</u>	Jun 03, 2013
<u>AB</u>	IMPAX LABS INC	<u>EQ 25MG HYDROCHLORIDE</u>	<u>A040156</u>	<u>001</u>	Jul 15, 1996
<u>AB</u>		<u>EQ 50MG HYDROCHLORIDE</u>	<u>A040156</u>	<u>002</u>	Jul 15, 1996
<u>AB</u>	SANDOZ	<u>EQ 25MG HYDROCHLORIDE</u>	<u>A087479</u>	<u>001</u>	
<u>AB</u>		<u>EQ 50MG HYDROCHLORIDE</u>	<u>A086183</u>	<u>001</u>	
<u>VISTARIL</u>					
<u>AB</u>	+ PFIZER	<u>EQ 25MG HYDROCHLORIDE</u>	<u>N011459</u>	<u>002</u>	
<u>AB</u>	+	<u>EQ 50MG HYDROCHLORIDE</u>	<u>N011459</u>	<u>004</u>	
HYDROXYZINE PAMOATE					
	BARR	<u>EQ 100MG HYDROCHLORIDE</u>	<u>A088488</u>	<u>001</u>	Jun 15, 1984

IBANDRONATE SODIUM

INJECTABLE; INTRAVENOUS

IBANDRONATE SODIUM

<u>AP</u>	ACCORD HLTHCARE	<u>EQ 3MG BASE/3ML</u>	<u>A206058</u>	<u>001</u>	Feb 05, 2016
<u>AP</u>	!	<u>EQ 3MG BASE/3ML</u>	<u>A204222</u>	<u>001</u>	Oct 16, 2015
<u>AP</u>	CHEMI SPA	<u>EQ 3MG BASE/3ML</u>	<u>A202235</u>	<u>001</u>	Sep 02, 2014
<u>AP</u>	EUGIA PHARMA	<u>EQ 3MG BASE/3ML</u>	<u>A205332</u>	<u>001</u>	Aug 19, 2015
<u>AP</u>	MYLAN LABS LTD	<u>EQ 3MG BASE/3ML</u>	<u>A202671</u>	<u>001</u>	Sep 02, 2014

TABLET; ORAL

IBANDRONATE SODIUM

<u>AB</u>	APOTEX INC	<u>EQ 150MG BASE</u>	<u>A078948</u>	<u>001</u>	Mar 19, 2012
<u>AB</u>	AUROBINDO PHARMA	<u>EQ 150MG BASE</u>	<u>A204502</u>	<u>001</u>	Mar 11, 2016
<u>AB</u>	!	<u>EQ 150MG BASE</u>	<u>A078997</u>	<u>001</u>	Apr 30, 2012
<u>AB</u>	MACLEODS PHARMS LTD	<u>EQ 150MG BASE</u>	<u>A206887</u>	<u>001</u>	Oct 31, 2017
<u>AB</u>	ORBION PHARMS	<u>EQ 150MG BASE</u>	<u>A078998</u>	<u>001</u>	Mar 19, 2012

PRESCRIPTION DRUG PRODUCT LIST

IBREXAFUNGERP CITRATE

TABLET; ORAL

BREXAFEMME

+! SCYNEXIS EQ 150MG BASE N214900 001 Jun 01, 2021

IBRUTINIB

CAPSULE; ORAL

IMBRUVICA

+ PHARMACYCLICS LLC 70MG N205552 002 Dec 20, 2017

+! 140MG N205552 001 Nov 13, 2013

SUSPENSION; ORAL

IMBRUVICA

+! PHARMACYCLICS LLC 70MG/ML N217003 001 Aug 24, 2022

TABLET; ORAL

IMBRUVICA

+ PHARMACYCLICS LLC 140MG N210563 001 Feb 16, 2018

+ 280MG N210563 002 Feb 16, 2018

+! 420MG N210563 003 Feb 16, 2018

IBUPROFEN

SOLUTION; INTRAVENOUS

CALDOLOR

+! CUMBERLAND PHARMS 800MG/8ML (100MG/ML) N022348 002 Jun 11, 2009

+! 800MG/200ML (4MG/ML) N022348 003 Jan 25, 2019

SUSPENSION; ORAL

IBUPROFEN

AB	!	ACTAVIS MID ATLANTIC	100MG/5ML	A074978 001	Mar 25, 1998
AB		AUROBINDO PHARMA LTD	100MG/5ML	A209178 001	Feb 16, 2018
AB		PADAGIS US	100MG/5ML	A076925 001	Sep 23, 2004
AB		STRIDES PHARMA	100MG/5ML	A215311 001	May 27, 2022
AB		TARO	100MG/5ML	A209204 001	Jun 23, 2017

TABLET; ORAL

IBUPROFEN

AB		ALKEM LABS LTD	400MG	A214699 001	Sep 13, 2021
AB			600MG	A214699 002	Sep 13, 2021
AB			800MG	A214699 003	Sep 13, 2021
AB		AMNEAL PHARMS NY	400MG	A071334 001	Nov 25, 1986
AB			400MG	A078558 001	Jun 18, 2007
AB			600MG	A071335 001	Nov 25, 1986
AB			600MG	A078558 002	Jun 18, 2007
AB			800MG	A071935 001	Oct 13, 1987
AB			800MG	A078558 003	Jun 18, 2007
AB		AUROBINDO PHARMA LTD	400MG	A213794 001	May 08, 2020
AB			600MG	A213794 002	May 08, 2020
AB			800MG	A213794 003	May 08, 2020
AB		CONTRACT PHARMACAL	400MG	A071268 002	Oct 15, 1986
AB			600MG	A071268 001	Oct 15, 1986
AB			800MG	A071268 003	Jul 01, 1988
AB		DR REDDYS	400MG	A075682 001	Nov 14, 2001
AB			600MG	A075682 002	Nov 14, 2001
AB	!		800MG	A075682 003	Nov 14, 2001
AB		DR REDDYS LABS INC	400MG	A076112 001	Oct 31, 2001
AB			600MG	A076112 002	Oct 31, 2001
AB			800MG	A076112 003	Oct 31, 2001
AB		GRANULES	400MG	A091625 001	Sep 15, 2015
AB			600MG	A091625 002	Sep 15, 2015
AB			800MG	A091625 003	Sep 15, 2015
AB		MARKSANS PHARMA	400MG	A090796 001	Dec 21, 2010
AB			600MG	A090796 002	Dec 21, 2010
AB			800MG	A090796 003	Dec 21, 2010
AB		SHANDONG XINHUA	400MG	A202413 001	Nov 23, 2016
AB			600MG	A202413 002	Nov 23, 2016
AB			800MG	A202413 003	Nov 23, 2016
AB		STRIDES PHARMA	400MG	A078329 001	Feb 05, 2009
AB			600MG	A078329 002	Feb 05, 2009
AB			800MG	A078329 003	Feb 05, 2009
AB		YICHANG HUMANWELL	400MG	A215318 001	Mar 30, 2022
AB			600MG	A215318 002	Mar 30, 2022
AB			800MG	A215318 003	Mar 30, 2022
AB		SHANDONG XINHUA	300MG	A202413 004	Jul 05, 2024

PRESCRIPTION DRUG PRODUCT LIST

IBUPROFEN LYSINE

INJECTABLE; INTRAVENOUS

IBUPROFEN LYSINE

AP	XGEN PHARMS	<u>EQ 20MG BASE/2ML (EQ 10MG BASE/ML)</u>	<u>A202402 001</u>	Mar 30, 2016
-----------	-------------	---	--------------------	--------------

NEOPROFEN

AP	+! RECORDATI RARE	<u>EQ 20MG BASE/2ML (EQ 10MG BASE/ML)</u>	<u>N021903 001</u>	Apr 13, 2006
-----------	-------------------	---	--------------------	--------------

IBUTILIDE FUMARATE

INJECTABLE; INJECTION

CORVERT

AP	+! PFIZER	<u>0.1MG/ML</u>	<u>N020491 001</u>	Dec 28, 1995
-----------	-----------	-----------------	--------------------	--------------

IBUTILIDE FUMARATE

AP	AVET LIFESCIENCES	<u>0.1MG/ML</u>	<u>A204146 001</u>	Apr 01, 2024
-----------	-------------------	-----------------	--------------------	--------------

AP	MYLAN INSTITUTIONAL	<u>0.1MG/ML</u>	<u>A090643 001</u>	Jan 11, 2010
-----------	---------------------	-----------------	--------------------	--------------

ICATIBANT ACETATE

INJECTABLE; SUBCUTANEOUS

FIRAZYR

AP	+! TAKEDA PHARMS USA	<u>EQ 30MG BASE/3ML (EQ 10MG BASE/ML)</u>	<u>N022150 001</u>	Aug 25, 2011
-----------	----------------------	---	--------------------	--------------

ICATIBANT ACETATE

AP	ALEMBIC	<u>EQ 30MG BASE/3ML (EQ 10MG BASE/ML)</u>	<u>A213773 001</u>	Jun 14, 2024
-----------	---------	---	--------------------	--------------

AP	CIPLA	<u>EQ 30MG BASE/3ML (EQ 10MG BASE/ML)</u>	<u>A212446 001</u>	Jul 13, 2020
-----------	-------	---	--------------------	--------------

AP	EUGIA PHARMA	<u>EQ 30MG BASE/3ML (EQ 10MG BASE/ML)</u>	<u>A213521 001</u>	Aug 14, 2023
-----------	--------------	---	--------------------	--------------

AP	FRESENIUS KABI USA	<u>EQ 30MG BASE/3ML (EQ 10MG BASE/ML)</u>	<u>A208317 001</u>	Jun 18, 2020
-----------	--------------------	---	--------------------	--------------

AP	JIANGSU HANSOH PHARM	<u>EQ 30MG BASE/3ML (EQ 10MG BASE/ML)</u>	<u>A211021 001</u>	Mar 09, 2020
-----------	----------------------	---	--------------------	--------------

AP	NANG KUANG PHARM CO	<u>EQ 30MG BASE/3ML (EQ 10MG BASE/ML)</u>	<u>A212081 001</u>	Dec 16, 2020
-----------	---------------------	---	--------------------	--------------

AP	TEVA PHARMS USA	<u>EQ 30MG BASE/3ML (EQ 10MG BASE/ML)</u>	<u>A210118 001</u>	Jul 15, 2019
-----------	-----------------	---	--------------------	--------------

AP	WILSHIRE PHARMS INC	<u>EQ 30MG BASE/3ML (EQ 10MG BASE/ML)</u>	<u>A211501 001</u>	Sep 01, 2020
-----------	---------------------	---	--------------------	--------------

ICODEXTRIN

SOLUTION; INTRAPERITONEAL

EXTRANEAL

+!	BAXTER HLTHCARE	7.5GM/100ML	N021321 001	Dec 20, 2002
----	-----------------	-------------	-------------	--------------

ICOSAPENT ETHYL

CAPSULE; ORAL

ICOSAPENT ETHYL

AB	APOTEX	<u>1GM</u>	<u>A209437 001</u>	Jun 30, 2021
-----------	--------	------------	--------------------	--------------

AB	ASCENT PHARMS INC	<u>500MG</u>	<u>A216811 001</u>	Dec 07, 2023
-----------	-------------------	--------------	--------------------	--------------

AB		<u>1GM</u>	<u>A216811 002</u>	Feb 16, 2024
-----------	--	------------	--------------------	--------------

AB	DR REDDYS	<u>500MG</u>	<u>A209499 002</u>	Mar 08, 2023
-----------	-----------	--------------	--------------------	--------------

AB		<u>1GM</u>	<u>A209499 001</u>	Aug 07, 2020
-----------	--	------------	--------------------	--------------

AB	HIKMA	<u>500MG</u>	<u>A209457 002</u>	Mar 08, 2023
-----------	-------	--------------	--------------------	--------------

AB		<u>1GM</u>	<u>A209457 001</u>	May 21, 2020
-----------	--	------------	--------------------	--------------

AB	HUMANWELL PURACAP	<u>500MG</u>	<u>A217919 001</u>	Dec 22, 2023
-----------	-------------------	--------------	--------------------	--------------

AB		<u>1GM</u>	<u>A217919 002</u>	Dec 22, 2023
-----------	--	------------	--------------------	--------------

AB	QILU	<u>1GM</u>	<u>A218899 001</u>	Nov 20, 2024
-----------	------	------------	--------------------	--------------

AB	SPRIASO LLC	<u>1GM</u>	<u>A218994 001</u>	Dec 20, 2024
-----------	-------------	------------	--------------------	--------------

AB	STRIDES SOFTGELS	<u>500MG</u>	<u>A217844 001</u>	Sep 22, 2023
-----------	------------------	--------------	--------------------	--------------

AB		<u>1GM</u>	<u>A217844 002</u>	Sep 22, 2023
-----------	--	------------	--------------------	--------------

AB	TEVA PHARMS USA	<u>500MG</u>	<u>A209525 001</u>	Sep 11, 2020
-----------	-----------------	--------------	--------------------	--------------

AB		<u>1GM</u>	<u>A209525 002</u>	Sep 11, 2020
-----------	--	------------	--------------------	--------------

AB	ZYDUS LIFESCIENCES	<u>500MG</u>	<u>A217656 001</u>	Apr 20, 2023
-----------	--------------------	--------------	--------------------	--------------

AB		<u>1GM</u>	<u>A217656 002</u>	Apr 20, 2023
-----------	--	------------	--------------------	--------------

VASCEPA

AB	+ AMARIN PHARMS	<u>500MG</u>	<u>N202057 002</u>	Feb 16, 2017
-----------	-----------------	--------------	--------------------	--------------

AB	+!	<u>1GM</u>	<u>N202057 001</u>	Jul 26, 2012
-----------	----	------------	--------------------	--------------

IDARUBICIN HYDROCHLORIDE

INJECTABLE; INJECTION

IDAMYCIN PFS

AP	+! PFIZER	<u>1MG/ML</u>	<u>N050734 001</u>	Feb 17, 1997
-----------	-----------	---------------	--------------------	--------------

IDARUBICIN HYDROCHLORIDE

AP	HIKMA	<u>1MG/ML</u>	<u>A065275 001</u>	Dec 14, 2006
-----------	-------	---------------	--------------------	--------------

AP		<u>1MG/ML</u>	<u>A065288 001</u>	May 15, 2007
-----------	--	---------------	--------------------	--------------

AP	MEITHEAL	<u>1MG/ML</u>	<u>A065036 001</u>	May 01, 2002
-----------	----------	---------------	--------------------	--------------

IDELALISIB

TABLET; ORAL

ZYDELIG

+	GILEAD SCIENCES INC	100MG	N205858 001	Jul 23, 2014
---	---------------------	-------	-------------	--------------

+!		150MG	N205858 002	Jul 23, 2014
----	--	-------	-------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

IFOSFAMIDE

INJECTABLE; INJECTION

IFEX

<u>AP</u>	+	BAXTER HLTHCARE	<u>1GM/VIAL</u>	<u>N019763 001</u>	Dec 30, 1988
<u>AP</u>	+		<u>3GM/VIAL</u>	<u>N019763 002</u>	Dec 30, 1988

IFOSFAMIDE

<u>AP</u>	!	FRESENIUS KABI USA	<u>1GM/VIAL</u>	<u>A076078 001</u>	May 28, 2002
<u>AP</u>	!		<u>3GM/VIAL</u>	<u>A076078 002</u>	May 28, 2002
<u>AP</u>		HIKMA	<u>1GM/20ML (50MG/ML)</u>	<u>A076619 001</u>	Jun 29, 2011
<u>AP</u>			<u>3GM/60ML (50MG/ML)</u>	<u>A076619 002</u>	Jun 29, 2011
<u>AP</u>	!	MEITHEAL	<u>1GM/20ML (50MG/ML)</u>	<u>A076657 001</u>	Apr 04, 2007
<u>AP</u>	!		<u>3GM/60ML (50MG/ML)</u>	<u>A076657 002</u>	Apr 04, 2007

ILOPERIDONE

TABLET; ORAL

FANAPT

+	!	VANDA PHARMS INC	1MG	N022192 001	May 06, 2009
	+		2MG	N022192 002	May 06, 2009
	+		4MG	N022192 003	May 06, 2009
	+		6MG	N022192 004	May 06, 2009
	+		8MG	N022192 005	May 06, 2009
	+		10MG	N022192 006	May 06, 2009
	+		12MG	N022192 007	May 06, 2009

ILOPROST

SOLUTION; INTRAVENOUS

AURLUMYN

+	!	BTG INTL	100MCG/ML (100MCG/ML)	N217933 001	Feb 13, 2024
---	---	----------	-----------------------	-------------	--------------

IMATINIB MESYLATE

SOLUTION; ORAL

IMKELDI

+	!	SHORLA ONCOLOGY	EQ 80MG BASE/ML	N219097 001	Nov 22, 2024
---	---	-----------------	-----------------	-------------	--------------

TABLET; ORAL

GLEEVEC

<u>AB</u>	+	NOVARTIS	<u>EQ 100MG BASE</u>	<u>N021588 001</u>	Apr 18, 2003
<u>AB</u>	+		<u>EQ 400MG BASE</u>	<u>N021588 002</u>	Apr 18, 2003

IMATINIB MESYLATE

<u>AB</u>		APOTEX	<u>EQ 100MG BASE</u>	<u>A079179 001</u>	Aug 05, 2016
<u>AB</u>			<u>EQ 400MG BASE</u>	<u>A079179 002</u>	Aug 05, 2016
<u>AB</u>		CHARTWELL RX	<u>EQ 100MG BASE</u>	<u>A208429 001</u>	Jan 17, 2019
<u>AB</u>			<u>EQ 400MG BASE</u>	<u>A208429 002</u>	Jan 17, 2019
<u>AB</u>		DR REDDYS	<u>EQ 100MG BASE</u>	<u>A206547 001</u>	Aug 13, 2018
<u>AB</u>			<u>EQ 400MG BASE</u>	<u>A206547 002</u>	Aug 13, 2018
<u>AB</u>		EUGIA PHARMA	<u>EQ 100MG BASE</u>	<u>A212773 001</u>	Jul 23, 2020
<u>AB</u>			<u>EQ 400MG BASE</u>	<u>A212773 002</u>	Jul 23, 2020
<u>AB</u>		MYLAN	<u>EQ 100MG BASE</u>	<u>A204644 001</u>	Jun 21, 2017
<u>AB</u>			<u>EQ 400MG BASE</u>	<u>A204644 002</u>	Jun 21, 2017
<u>AB</u>		NATCO PHARMA LTD	<u>EQ 100MG BASE</u>	<u>A207818 001</u>	Mar 01, 2019
<u>AB</u>			<u>EQ 400MG BASE</u>	<u>A207818 002</u>	Mar 01, 2019
<u>AB</u>		QILU PHARM HAINAN	<u>EQ 100MG BASE</u>	<u>A212135 001</u>	Jun 21, 2022
<u>AB</u>			<u>EQ 400MG BASE</u>	<u>A212135 002</u>	Jun 21, 2022
<u>AB</u>		SHILPA	<u>EQ 100MG BASE</u>	<u>A208302 001</u>	Jan 17, 2019
<u>AB</u>			<u>EQ 400MG BASE</u>	<u>A208302 002</u>	Jan 17, 2019
<u>AB</u>		SUN PHARM	<u>EQ 100MG BASE</u>	<u>A078340 001</u>	Dec 03, 2015
<u>AB</u>			<u>EQ 400MG BASE</u>	<u>A078340 002</u>	Dec 03, 2015
<u>AB</u>		TEVA PHARMS USA	<u>EQ 100MG BASE</u>	<u>A204285 001</u>	Aug 04, 2016
<u>AB</u>			<u>EQ 400MG BASE</u>	<u>A204285 002</u>	Aug 04, 2016
<u>AB</u>		ZYDUS PHARMS	<u>EQ 100MG BASE</u>	<u>A210658 001</u>	Apr 08, 2020
<u>AB</u>			<u>EQ 400MG BASE</u>	<u>A210658 002</u>	Apr 08, 2020

IMETELSTAT SODIUM

POWDER; INTRAVENOUS

RYTELO

+	!	GERON	EQ 47MG BASE/VIAL	N217779 001	Jun 06, 2024
+	!		EQ 188MG BASE/VIAL	N217779 002	Jun 06, 2024

IMIPRAMINE HYDROCHLORIDE

TABLET; ORAL

IMIPRAMINE HYDROCHLORIDE

<u>AB</u>		LEADING	<u>10MG</u>	<u>A040903 001</u>	Oct 24, 2012
<u>AB</u>			<u>25MG</u>	<u>A040903 002</u>	Oct 24, 2012
<u>AB</u>	!		<u>50MG</u>	<u>A040903 003</u>	Oct 24, 2012
<u>AB</u>		OXFORD PHARMS	<u>10MG</u>	<u>A040751 003</u>	Feb 28, 2008
<u>AB</u>			<u>25MG</u>	<u>A040751 002</u>	Feb 28, 2008

PRESCRIPTION DRUG PRODUCT LIST

IMIPRAMINE HYDROCHLORIDE

TABLET; ORAL

IMIPRAMINE HYDROCHLORIDE

<u>AB</u>		<u>50MG</u>	<u>A040751 001</u>	Feb 28, 2008
<u>AB</u>	SANDOZ	<u>10MG</u>	<u>A084936 002</u>	
<u>AB</u>		<u>25MG</u>	<u>A083745 001</u>	
<u>AB</u>		<u>50MG</u>	<u>A084937 001</u>	
<u>AB</u>	STRIDES PHARMA	<u>10MG</u>	<u>A088292 001</u>	Oct 21, 1983
<u>AB</u>		<u>25MG</u>	<u>A088262 001</u>	Oct 21, 1983
<u>AB</u>		<u>50MG</u>	<u>A088276 001</u>	Oct 21, 1983
<u>AB</u>	SUN PHARM INDUSTRIES	<u>25MG</u>	<u>A081049 001</u>	Jun 05, 1990
<u>AB</u>		<u>50MG</u>	<u>A081050 001</u>	Jun 05, 1990

IMIPRAMINE PAMOATE

CAPSULE; ORAL

IMIPRAMINE PAMOATE

<u>AB</u>	!	HIKMA	<u>EQ 75MG HYDROCHLORIDE</u>	<u>A091099 001</u>	Apr 16, 2010
<u>AB</u>			<u>EQ 100MG HYDROCHLORIDE</u>	<u>A091099 002</u>	Apr 16, 2010
<u>AB</u>			<u>EQ 125MG HYDROCHLORIDE</u>	<u>A091099 003</u>	Apr 16, 2010
<u>AB</u>			<u>EQ 150MG HYDROCHLORIDE</u>	<u>A091099 004</u>	Apr 16, 2010
<u>AB</u>		LUPIN LTD	<u>EQ 75MG HYDROCHLORIDE</u>	<u>A090444 001</u>	Apr 16, 2010
<u>AB</u>			<u>EQ 100MG HYDROCHLORIDE</u>	<u>A090444 002</u>	Apr 16, 2010
<u>AB</u>			<u>EQ 125MG HYDROCHLORIDE</u>	<u>A090444 003</u>	Apr 16, 2010
<u>AB</u>			<u>EQ 150MG HYDROCHLORIDE</u>	<u>A090444 004</u>	Apr 16, 2010

IMIQUIMOD

CREAM; TOPICAL

IMIQUIMOD

<u>AB</u>		APOTEX INC	<u>5%</u>	<u>A091308 001</u>	Apr 06, 2012
<u>AB</u>		FOUGERA PHARMS	<u>5%</u>	<u>A078548 001</u>	Feb 25, 2010
<u>AB</u>	!	GLENMARK PHARMS INC	<u>5%</u>	<u>A201994 001</u>	Mar 06, 2012
<u>AB</u>		PADAGIS ISRAEL	<u>5%</u>	<u>A078837 001</u>	Sep 07, 2010
<u>AB</u>		TARO	<u>3.75%</u>	<u>A205971 001</u>	Jan 26, 2021
<u>AB</u>			<u>5%</u>	<u>A200173 001</u>	Apr 15, 2011

ZYCLARA

<u>AB</u>	+	BAUSCH	<u>3.75%</u>	<u>N022483 001</u>	Mar 25, 2010
	+		<u>2.5%</u>	<u>N022483 002</u>	Jul 15, 2011

INAMRINONE LACTATE

INJECTABLE; INJECTION

AMRINONE LACTATE

!	HIKMA	EQ 5MG BASE/ML	A075513 001	May 09, 2000
---	-------	----------------	-------------	--------------

INAVOLISIB

TABLET; ORAL

ITOVEBI

+	GENENTECH INC	3MG	N219249 001	Oct 10, 2024
+	!	9MG	N219249 002	Oct 10, 2024

INCLISIRAN SODIUM

SOLUTION; SUBCUTANEOUS

LEQVIO

+	NOVARTIS	EQ 284MG BASE/1.5ML (EQ 189MG BASE/ML)	N214012 001	Dec 22, 2021
---	----------	--	-------------	--------------

INDAPAMIDE

TABLET; ORAL

INDAPAMIDE

<u>AB</u>		ACTAVIS ELIZABETH	<u>1.25MG</u>	<u>A074722 001</u>	Jun 17, 1996
<u>AB</u>			<u>2.5MG</u>	<u>A074722 002</u>	Jun 17, 1996
<u>AB</u>		ANI PHARMS	<u>1.25MG</u>	<u>A074299 002</u>	Apr 29, 1996
<u>AB</u>	!		<u>2.5MG</u>	<u>A074299 001</u>	Jul 27, 1995
<u>AB</u>		RISING	<u>1.25MG</u>	<u>A074461 002</u>	Mar 26, 1997
<u>AB</u>			<u>2.5MG</u>	<u>A074461 001</u>	Mar 27, 1996

INDIGOTINDISULFONATE SODIUM

SOLUTION; INTRAVENOUS

BLUDIGO

+	PROVEPHARM SAS	40MG/5ML (8MG/ML)	N216264 001	Jul 08, 2022
---	----------------	-------------------	-------------	--------------

INDIUM IN-111 OXYQUINOLINE

INJECTABLE; INJECTION

INDIUM IN 111 OXYQUINOLINE

<u>AP</u>		BWXT ITG	<u>1mCi/ML</u>	<u>A202586 001</u>	Jul 25, 2018
<u>AP</u>	+	GE HEALTHCARE	<u>1mCi/ML</u>	<u>N019044 001</u>	Dec 24, 1985

PRESCRIPTION DRUG PRODUCT LIST

INDIUM IN-111 PENTETATE DISODIUM

INJECTABLE; INTRATHECAL

MPI INDIUM DTPA IN 111

+! GE HEALTHCARE 1mCi/ML

N017707 001 Feb 18, 1982

INDIUM IN-111 PENTETREOTIDE KIT

INJECTABLE; INJECTION

INDIUM IN-111 PENTETREOTIDE KITAP SUN PHARM INDS INC 3mCi/MLA212785 001 Jul 01, 2024OCTREOSCANAP +! CURIUM 3mCi/MLN020314 001 Jun 02, 1994INDOCYANINE GREEN

INJECTABLE; INJECTION

INDOCYANINE GREEN

! RENEW PHARMS 25MG/VIAL

A040811 001 Nov 21, 2007

POWDER; INTRAVENOUS, INTERSTITIAL

SPY AGENT GREEN KIT

+! NOVADAQ TECH 25MG/VIAL

N211580 001 Nov 21, 2018

INDOMETHACIN

CAPSULE; ORAL

INDOMETHACINAB CHARTWELL MOLECULES 25MGN018829 002 Aug 06, 1984AB 50MGA070651 001 Mar 05, 1986AB 50MGN018829 001 Aug 06, 1984AB GLENMARK PHARMS LTD 25MGA091276 001 Dec 22, 2010AB ! 50MGA091276 002 Dec 22, 2010AB HETERO LABS LTD III 25MGA091240 001 Apr 12, 2011AB 50MGA091240 002 Apr 12, 2011AB SANDOZ 25MGA070673 001 Apr 29, 1987AB 50MGA070674 001 Apr 29, 1987AB ZYDUS LIFESCIENCES 25MGA090403 001 Nov 15, 2010AB 50MGA090403 002 Nov 15, 2010

CAPSULE, EXTENDED RELEASE; ORAL

INDOMETHACINAB AMNEAL PHARMS 75MGA091549 001 Dec 01, 2010AB AVANTHI INC 75MGA079175 001 Mar 06, 2009AB CHARTWELL RX 75MGA200529 001 Nov 30, 2010AB GLENMARK PHARMS LTD 75MGA203501 001 Jun 22, 2017AB ! HETERO LABS LTD III 75MGA201807 001 Sep 28, 2012AB NOVAST LABS 75MGA204853 001 May 08, 2017AB ZYDUS PHARMS 75MGA202711 001 Sep 25, 2017

INJECTABLE; INJECTION

INDOMETHACIN

+! FRESENIUS KABI USA EQ 1MG BASE/VIAL

N022536 001 Mar 17, 2010

SUPPOSITORY; RECTAL

INDOMETHACINAB ! COSETTE 50MGA073314 001 Aug 31, 1992AB HIKMA 50MGA215899 001 Nov 19, 2024AB ZYDUS LIFESCIENCES 50MGA216184 001 Aug 02, 2023

SUSPENSION; ORAL

INDOCINAB +! ZYLA 25MG/5MLN018332 001 Oct 10, 1985INDOMETHACINAB NOVITIUM PHARMA 25MG/5MLA217883 001 Jan 12, 2024IOBENGUANE SULFATE I-123

SOLUTION; INTRAVENOUS

ADREVIEWAP +! GE HEALTHCARE 10mCi/5ML (2mCi/ML)N022290 001 Sep 19, 2008IOBENGUANE I-123AP BWXT MEDCL 10mCi/5ML (2mCi/ML)A213637 001 Dec 09, 2024IODIXANOL

INJECTABLE; INJECTION

IODIXANOLAP HENGRUI PHARMA 55%A214271 001 May 19, 2022AP 65.2%A214271 002 May 19, 2022VISIPAQUE 270AP +! GE HEALTHCARE 55%N020351 001 Mar 22, 1996VISIPAQUE 320AP +! GE HEALTHCARE 65.2%N020351 002 Mar 22, 1996

65.2%

N020808 002 Aug 29, 1997

PRESCRIPTION DRUG PRODUCT LIST

IOFLUPANE I-123

SOLUTION; INTRAVENOUS

DATSCAN

AP	+!	GE HLTHCARE INC	<u>5mCi/2.5ML (2mCi/ML)</u>	<u>N022454 001</u>	Jan 14, 2011
-----------	-----------	-----------------	------------------------------------	---------------------------	--------------

IOFLUPANE I-123

AP		CURIUM	<u>5mCi/2.5ML (2mCi/ML)</u>	<u>A213792 001</u>	Mar 30, 2022
-----------	--	--------	------------------------------------	---------------------------	--------------

IOHEXOL

INJECTABLE; INJECTION

OMNIPAQUE 140

+!	GE HEALTHCARE	30.2%		N018956 005	Nov 30, 1988
-----------	---------------	-------	--	-------------	--------------

SOLUTION; INJECTION, ORAL

OMNIPAQUE 350

+!	GE HEALTHCARE	75.5%		N018956 004	Dec 26, 1985
-----------	---------------	-------	--	-------------	--------------

		75.5%		N020608 003	Oct 24, 1995
--	--	-------	--	-------------	--------------

SOLUTION; INJECTION, ORAL, RECTAL

OMNIPAQUE 180

+!	GE HEALTHCARE	38.8%		N018956 001	Dec 26, 1985
-----------	---------------	-------	--	-------------	--------------

OMNIPAQUE 240

+!	GE HEALTHCARE	51.8%		N018956 002	Dec 26, 1985
-----------	---------------	-------	--	-------------	--------------

OMNIPAQUE 300

+!	GE HEALTHCARE	64.7%		N018956 003	Dec 26, 1985
-----------	---------------	-------	--	-------------	--------------

		64.7%		N020608 002	Oct 24, 1995
--	--	-------	--	-------------	--------------

SOLUTION; ORAL

OMNIPAQUE 12

+!	GE HEALTHCARE	2.6%		N018956 009	Apr 17, 2018
-----------	---------------	------	--	-------------	--------------

OMNIPAQUE 9

+!	GE HEALTHCARE	1.9%		N018956 008	Apr 17, 2018
-----------	---------------	------	--	-------------	--------------

IOMEPROL

SOLUTION; INTRA-ARTERIAL

IOMERUVU

+!	BRACCO	15GM IODINE/50ML (300MG IODINE/ML)		N216016 002	Nov 27, 2024
-----------	--------	------------------------------------	--	-------------	--------------

+!		17.5GM IODINE/50ML (350MG IODINE/ML)		N216016 006	Nov 27, 2024
-----------	--	--------------------------------------	--	-------------	--------------

+!		20GM IODINE/50ML (400MG IODINE/ML)		N216016 010	Nov 27, 2024
-----------	--	------------------------------------	--	-------------	--------------

+!		25GM IODINE/100ML (250MG IODINE/ML)		N216016 001	Nov 27, 2024
-----------	--	-------------------------------------	--	-------------	--------------

+!		30GM IODINE/100ML (300MG IODINE/ML)		N216016 003	Nov 27, 2024
-----------	--	-------------------------------------	--	-------------	--------------

+!		35GM IODINE/100ML (350MG IODINE/ML)		N216016 007	Nov 27, 2024
-----------	--	-------------------------------------	--	-------------	--------------

+!		40GM IODINE/100ML (400MG IODINE/ML)		N216016 011	Nov 27, 2024
-----------	--	-------------------------------------	--	-------------	--------------

+!		60GM IODINE/150ML (400MG IODINE/ML)		N216016 012	Nov 27, 2024
-----------	--	-------------------------------------	--	-------------	--------------

+!		45GM IODINE/150ML (300MG IODINE/ML)		N216016 004	Nov 27, 2024
-----------	--	-------------------------------------	--	-------------	--------------

+!		52.5GM IODINE/150ML (350MG IODINE/ML)		N216016 008	Nov 27, 2024
-----------	--	---------------------------------------	--	-------------	--------------

+!		60GM IODINE/200ML (300MG IODINE/ML)		N216016 005	Nov 27, 2024
-----------	--	-------------------------------------	--	-------------	--------------

+!		70GM IODINE/200ML (350MG IODINE/ML)		N216016 009	Nov 27, 2024
-----------	--	-------------------------------------	--	-------------	--------------

+!		80GM IODINE/200ML (400MG IODINE/ML)		N216016 013	Nov 27, 2024
-----------	--	-------------------------------------	--	-------------	--------------

SOLUTION; INTRAVENOUS

IOMERUVU

+!	BRACCO	15GM IODINE/50ML (300MG IODINE/ML)		N216017 002	Nov 27, 2024
-----------	--------	------------------------------------	--	-------------	--------------

+!		17.5GM IODINE/50ML (350MG IODINE/ML)		N216017 006	Nov 27, 2024
-----------	--	--------------------------------------	--	-------------	--------------

+!		20GM IODINE/50ML (400MG IODINE/ML)		N216017 010	Nov 27, 2024
-----------	--	------------------------------------	--	-------------	--------------

+!		25GM IODINE/100ML (250MG IODINE/ML)		N216017 001	Nov 27, 2024
-----------	--	-------------------------------------	--	-------------	--------------

+!		30GM IODINE/100ML (300MG IODINE/ML)		N216017 003	Nov 27, 2024
-----------	--	-------------------------------------	--	-------------	--------------

+!		35GM IODINE/100ML (350MG IODINE/ML)		N216017 007	Nov 27, 2024
-----------	--	-------------------------------------	--	-------------	--------------

+!		40GM IODINE/100ML (400MG IODINE/ML)		N216017 011	Nov 27, 2024
-----------	--	-------------------------------------	--	-------------	--------------

+!		60GM IODINE/150ML (400MG IODINE/ML)		N216017 012	Nov 27, 2024
-----------	--	-------------------------------------	--	-------------	--------------

+!		45GM IODINE/150ML (300MG IODINE/ML)		N216017 004	Nov 27, 2024
-----------	--	-------------------------------------	--	-------------	--------------

+!		52.5GM IODINE/150ML (350MG IODINE/ML)		N216017 008	Nov 27, 2024
-----------	--	---------------------------------------	--	-------------	--------------

+!		60GM IODINE/200ML (300MG IODINE/ML)		N216017 005	Nov 27, 2024
-----------	--	-------------------------------------	--	-------------	--------------

+!		70GM IODINE/200ML (350MG IODINE/ML)		N216017 009	Nov 27, 2024
-----------	--	-------------------------------------	--	-------------	--------------

+!		80GM IODINE/200ML (400MG IODINE/ML)		N216017 013	Nov 27, 2024
-----------	--	-------------------------------------	--	-------------	--------------

IOPAMIDOL

INJECTABLE; INJECTION

IOPAMIDOL

AP		HAINAN POLY	<u>51%</u>	<u>A217134 002</u>	Apr 09, 2024
-----------	--	-------------	-------------------	---------------------------	--------------

AP			<u>61%</u>	<u>A215382 002</u>	Feb 27, 2023
-----------	--	--	-------------------	---------------------------	--------------

AP			<u>61%</u>	<u>A217134 003</u>	Apr 09, 2024
-----------	--	--	-------------------	---------------------------	--------------

AP			<u>76%</u>	<u>A217134 004</u>	Apr 09, 2024
-----------	--	--	-------------------	---------------------------	--------------

ISOVUE-250

AP	+!	BRACCO	<u>51%</u>	<u>N018735 007</u>	Jul 06, 1992
-----------	-----------	--------	-------------------	---------------------------	--------------

ISOVUE-300

AP	+!	BRACCO	<u>61%</u>	<u>N018735 002</u>	Dec 31, 1985
-----------	-----------	--------	-------------------	---------------------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

IOPAMIDOL

INJECTABLE; INJECTION

ISOVUE-370**AP** +! BRACCO **76%** **N018735 003** Dec 31, 1985ISOVUE-M 300**AP** +! BRACCO **61%** **N018735 004** Dec 31, 1985IOPAMIDOL**AP1** HAINAN POLY **41%** **A215382 001** Feb 27, 2023ISOVUE-M 200**AP1** +! BRACCO **41%** **N018735 001** Dec 31, 1985IOPAMIDOL**AP2** HAINAN POLY **41%** **A217134 001** Sep 27, 2023ISOVUE-200**AP2** +! BRACCO **41%** **N018735 006** Jul 07, 1987

ISOVUE-300

+! BRACCO 61% N020327 003 Oct 12, 1994

ISOVUE-370

+! BRACCO 76% N020327 004 Oct 12, 1994

IOPROMIDE

INJECTABLE; INJECTION

ULTRAVIST (PHARMACY BULK)

+! BAYER HLTHCARE 62.3% N021425 001 Sep 20, 2002

+! 76.9% N021425 002 Sep 20, 2002

ULTRAVIST 300

+! BAYER HLTHCARE 62.3% N020220 002 May 10, 1995

ULTRAVIST 370

+! BAYER HLTHCARE 76.9% N020220 001 May 10, 1995

IOTHALAMATE MEGLUMINE

INJECTABLE; INJECTION

CONRAY

+! LIEBEL-FLARSHEIM 60% N013295 001

SOLUTION; INTRAVESICAL

CYSTO-CONRAY II

LIEBEL-FLARSHEIM 17.2% N017057 002

IOTHALAMATE SODIUM I-125

INJECTABLE; INJECTION

GLOFIL-125

ISOTEX 250-300uCi/ML N017279 001

IOVERSOL

INJECTABLE; INJECTION

OPTIRAY 300

+! LIEBEL-FLARSHEIM 64% N019710 004 Jan 22, 1992

+! 64% N020923 004 May 13, 1999

OPTIRAY 320

+! LIEBEL-FLARSHEIM 68% N019710 001 Dec 30, 1988

+! 68% N020923 002 May 29, 1998

OPTIRAY 350

+! LIEBEL-FLARSHEIM 74% N019710 005 Jan 22, 1992

+! 74% N020923 003 May 28, 1998

IPRATROPIUM BROMIDE

AEROSOL, METERED; INHALATION

ATROVENT HFA

+! BOEHRINGER 0.021MG/INH N021527 001 Nov 27, 2004

INGELHEIM

SOLUTION; INHALATION

IPRATROPIUM BROMIDE**AN** LUOXIN AUROVITAS **0.02%** **A206543 001** Oct 27, 2016**AN** NEPHRON **0.02%** **A075562 001** Sep 27, 2001**AN** ! RITEDOSE CORP **0.02%** **A075693 001** Jan 26, 2001**AN** SUN PHARM **0.02%** **A207903 001** Jan 03, 2017

SPRAY, METERED; NASAL

IPRATROPIUM BROMIDE**AB** AMNEAL **0.021MG/SPRAY** **A215104 001** Aug 16, 2022**AB** **0.042MG/SPRAY** **A215105 001** Aug 16, 2022**AB** APOTEX INC **0.021MG/SPRAY** **A076156 001** Apr 18, 2003**AB** **0.042MG/SPRAY** **A076155 001** Apr 18, 2003**AB** BAUSCH **0.021MG/SPRAY** **A076025 001** Mar 31, 2003**AB** **0.042MG/SPRAY** **A076103 001** Mar 31, 2003**AB** ! HIKMA **0.021MG/SPRAY** **A076664 001** Nov 05, 2003**AB** ! **0.042MG/SPRAY** **A076598 001** Nov 05, 2003

PRESCRIPTION DRUG PRODUCT LIST

IPRATROPIUM BROMIDE

SPRAY, METERED;NASAL

IPRATROPIUM BROMIDE

<u>AB</u>	RUBICON	<u>0.042MG/SPRAY</u>	<u>A219221</u>	<u>001</u>	Nov 15, 2024
-----------	---------	----------------------	----------------	------------	--------------

IPTACOPAN HYDROCHLORIDE

CAPSULE;ORAL

FABHALTA

+! NOVARTIS

EQ 200MG BASE

N218276 001 Dec 05, 2023

IRBESARTAN

TABLET;ORAL

AVAPRO

<u>AB</u>	+ SANOFI AVENTIS US	<u>150MG</u>	<u>N020757</u>	<u>002</u>	Sep 30, 1997
-----------	---------------------	--------------	----------------	------------	--------------

<u>AB</u>	+!	<u>300MG</u>	<u>N020757</u>	<u>003</u>	Sep 30, 1997
-----------	----	--------------	----------------	------------	--------------

IRBESARTAN

<u>AB</u>	ALEMBIC PHARMS LTD	<u>75MG</u>	<u>A091236</u>	<u>001</u>	Oct 15, 2012
-----------	--------------------	-------------	----------------	------------	--------------

<u>AB</u>		<u>150MG</u>	<u>A091236</u>	<u>002</u>	Oct 15, 2012
-----------	--	--------------	----------------	------------	--------------

<u>AB</u>		<u>300MG</u>	<u>A091236</u>	<u>003</u>	Oct 15, 2012
-----------	--	--------------	----------------	------------	--------------

<u>AB</u>	AMNEAL PHARMS	<u>75MG</u>	<u>A204740</u>	<u>001</u>	Apr 17, 2018
-----------	---------------	-------------	----------------	------------	--------------

<u>AB</u>		<u>150MG</u>	<u>A204740</u>	<u>002</u>	Apr 17, 2018
-----------	--	--------------	----------------	------------	--------------

<u>AB</u>		<u>300MG</u>	<u>A204740</u>	<u>003</u>	Apr 17, 2018
-----------	--	--------------	----------------	------------	--------------

<u>AB</u>	AUROBINDO PHARMA	<u>75MG</u>	<u>A203081</u>	<u>001</u>	Sep 27, 2012
-----------	------------------	-------------	----------------	------------	--------------

<u>AB</u>	LTD				
-----------	-----	--	--	--	--

<u>AB</u>		<u>150MG</u>	<u>A203081</u>	<u>002</u>	Sep 27, 2012
-----------	--	--------------	----------------	------------	--------------

<u>AB</u>		<u>300MG</u>	<u>A203081</u>	<u>003</u>	Sep 27, 2012
-----------	--	--------------	----------------	------------	--------------

<u>AB</u>	CHARTWELL MOLECULAR	<u>75MG</u>	<u>A077205</u>	<u>001</u>	Nov 14, 2012
-----------	---------------------	-------------	----------------	------------	--------------

<u>AB</u>		<u>150MG</u>	<u>A077205</u>	<u>002</u>	Nov 14, 2012
-----------	--	--------------	----------------	------------	--------------

<u>AB</u>		<u>300MG</u>	<u>A077205</u>	<u>003</u>	Nov 14, 2012
-----------	--	--------------	----------------	------------	--------------

<u>AB</u>	HETERO LABS LTD V	<u>75MG</u>	<u>A202910</u>	<u>001</u>	Sep 27, 2012
-----------	-------------------	-------------	----------------	------------	--------------

<u>AB</u>		<u>150MG</u>	<u>A202910</u>	<u>002</u>	Sep 27, 2012
-----------	--	--------------	----------------	------------	--------------

<u>AB</u>		<u>300MG</u>	<u>A202910</u>	<u>003</u>	Sep 27, 2012
-----------	--	--------------	----------------	------------	--------------

<u>AB</u>	HISUN PHARM	<u>75MG</u>	<u>A206194</u>	<u>001</u>	Jun 14, 2016
-----------	-------------	-------------	----------------	------------	--------------

<u>AB</u>	HANGZHOU				
-----------	----------	--	--	--	--

<u>AB</u>		<u>150MG</u>	<u>A206194</u>	<u>002</u>	Jun 14, 2016
-----------	--	--------------	----------------	------------	--------------

<u>AB</u>		<u>300MG</u>	<u>A206194</u>	<u>003</u>	Jun 14, 2016
-----------	--	--------------	----------------	------------	--------------

<u>AB</u>	JUBILANT GENERICS	<u>75MG</u>	<u>A203534</u>	<u>001</u>	Feb 23, 2015
-----------	-------------------	-------------	----------------	------------	--------------

<u>AB</u>		<u>150MG</u>	<u>A203534</u>	<u>002</u>	Feb 23, 2015
-----------	--	--------------	----------------	------------	--------------

<u>AB</u>		<u>300MG</u>	<u>A203534</u>	<u>003</u>	Feb 23, 2015
-----------	--	--------------	----------------	------------	--------------

<u>AB</u>	LUPIN LTD	<u>75MG</u>	<u>A201531</u>	<u>001</u>	Oct 15, 2012
-----------	-----------	-------------	----------------	------------	--------------

<u>AB</u>		<u>150MG</u>	<u>A201531</u>	<u>002</u>	Oct 15, 2012
-----------	--	--------------	----------------	------------	--------------

<u>AB</u>		<u>300MG</u>	<u>A201531</u>	<u>003</u>	Oct 15, 2012
-----------	--	--------------	----------------	------------	--------------

<u>AB</u>	MACLEODS PHARMS LTD	<u>75MG</u>	<u>A202254</u>	<u>001</u>	Oct 03, 2012
-----------	---------------------	-------------	----------------	------------	--------------

<u>AB</u>		<u>150MG</u>	<u>A202254</u>	<u>002</u>	Oct 03, 2012
-----------	--	--------------	----------------	------------	--------------

<u>AB</u>		<u>300MG</u>	<u>A202254</u>	<u>003</u>	Oct 03, 2012
-----------	--	--------------	----------------	------------	--------------

<u>AB</u>	PRINSTON INC	<u>75MG</u>	<u>A203071</u>	<u>001</u>	Sep 27, 2012
-----------	--------------	-------------	----------------	------------	--------------

<u>AB</u>		<u>150MG</u>	<u>A203071</u>	<u>002</u>	Sep 27, 2012
-----------	--	--------------	----------------	------------	--------------

<u>AB</u>		<u>300MG</u>	<u>A203071</u>	<u>003</u>	Sep 27, 2012
-----------	--	--------------	----------------	------------	--------------

<u>AB</u>	SANDOZ	<u>75MG</u>	<u>A077466</u>	<u>001</u>	Sep 27, 2012
-----------	--------	-------------	----------------	------------	--------------

<u>AB</u>		<u>150MG</u>	<u>A077466</u>	<u>002</u>	Sep 27, 2012
-----------	--	--------------	----------------	------------	--------------

<u>AB</u>		<u>300MG</u>	<u>A077466</u>	<u>003</u>	Sep 27, 2012
-----------	--	--------------	----------------	------------	--------------

<u>AB</u>	SCIEGEN PHARMS INC	<u>75MG</u>	<u>A204774</u>	<u>001</u>	Dec 07, 2015
-----------	--------------------	-------------	----------------	------------	--------------

<u>AB</u>		<u>150MG</u>	<u>A204774</u>	<u>002</u>	Dec 07, 2015
-----------	--	--------------	----------------	------------	--------------

<u>AB</u>		<u>300MG</u>	<u>A204774</u>	<u>003</u>	Dec 07, 2015
-----------	--	--------------	----------------	------------	--------------

<u>AB</u>	TEVA PHARMS	<u>75MG</u>	<u>A077159</u>	<u>001</u>	Mar 30, 2012
-----------	-------------	-------------	----------------	------------	--------------

<u>AB</u>		<u>150MG</u>	<u>A077159</u>	<u>002</u>	Mar 30, 2012
-----------	--	--------------	----------------	------------	--------------

<u>AB</u>		<u>300MG</u>	<u>A077159</u>	<u>003</u>	Mar 30, 2012
-----------	--	--------------	----------------	------------	--------------

<u>AB</u>	UNICHEM	<u>75MG</u>	<u>A203020</u>	<u>001</u>	Dec 07, 2015
-----------	---------	-------------	----------------	------------	--------------

<u>AB</u>		<u>150MG</u>	<u>A203020</u>	<u>002</u>	Dec 07, 2015
-----------	--	--------------	----------------	------------	--------------

<u>AB</u>		<u>300MG</u>	<u>A203020</u>	<u>003</u>	Dec 07, 2015
-----------	--	--------------	----------------	------------	--------------

<u>AB</u>	ZYDUS PHARMS USA	<u>75MG</u>	<u>A079213</u>	<u>001</u>	Sep 27, 2012
-----------	------------------	-------------	----------------	------------	--------------

<u>AB</u>	INC				
-----------	-----	--	--	--	--

<u>AB</u>		<u>150MG</u>	<u>A079213</u>	<u>002</u>	Sep 27, 2012
-----------	--	--------------	----------------	------------	--------------

<u>AB</u>		<u>300MG</u>	<u>A079213</u>	<u>003</u>	Sep 27, 2012
-----------	--	--------------	----------------	------------	--------------

IRINOTECAN HYDROCHLORIDE

INJECTABLE; INJECTION

CAMPTOSAR

<u>AP</u>	+! PFIZER INC	<u>40MG/2ML (20MG/ML)</u>	<u>N020571</u>	<u>001</u>	Jun 14, 1996
-----------	---------------	---------------------------	----------------	------------	--------------

<u>AP</u>	+!	<u>100MG/5ML (20MG/ML)</u>	<u>N020571</u>	<u>002</u>	Jun 14, 1996
-----------	----	----------------------------	----------------	------------	--------------

<u>AP</u>	+!	<u>300MG/15ML (20MG/ML)</u>	<u>N020571</u>	<u>003</u>	Aug 05, 2010
-----------	----	-----------------------------	----------------	------------	--------------

IRINOTECAN HYDROCHLORIDE

<u>AP</u>	ACCORD HLTHCARE	<u>40MG/2ML (20MG/ML)</u>	<u>A079068</u>	<u>001</u>	Nov 21, 2008
-----------	-----------------	---------------------------	----------------	------------	--------------

<u>AP</u>		<u>100MG/5ML (20MG/ML)</u>	<u>A079068</u>	<u>002</u>	Nov 21, 2008
-----------	--	----------------------------	----------------	------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

IRINOTECAN HYDROCHLORIDE

INJECTABLE; INJECTION

IRINOTECAN HYDROCHLORIDE

<u>AP</u>	ACTAVIS TOTOWA	<u>40MG/2ML (20MG/ML)</u>	<u>A078589 001</u>	Feb 27, 2008
<u>AP</u>		<u>100MG/5ML (20MG/ML)</u>	<u>A078589 002</u>	Feb 27, 2008
<u>AP</u>		<u>500MG/25ML (20MG/ML)</u>	<u>A078589 003</u>	Nov 18, 2015
<u>AP</u>	EPIC PHARMA LLC	<u>40MG/2ML (20MG/ML)</u>	<u>A090726 001</u>	Sep 16, 2009
<u>AP</u>		<u>100MG/5ML (20MG/ML)</u>	<u>A090726 002</u>	Sep 16, 2009
<u>AP</u>	EUGIA PHARMA	<u>40MG/2ML (20MG/ML)</u>	<u>A213278 001</u>	Nov 02, 2020
<u>AP</u>		<u>100MG/5ML (20MG/ML)</u>	<u>A213278 002</u>	Nov 02, 2020
<u>AP</u>		<u>300MG/15ML (20MG/ML)</u>	<u>A213278 003</u>	Nov 02, 2020
<u>AP</u>		<u>500MG/25ML (20MG/ML)</u>	<u>A213278 004</u>	Apr 11, 2022
<u>AP</u>	FRESENIUS KABI USA	<u>40MG/2ML (20MG/ML)</u>	<u>A077776 001</u>	Feb 27, 2008
<u>AP</u>		<u>100MG/5ML (20MG/ML)</u>	<u>A077776 002</u>	Feb 27, 2008
<u>AP</u>	GLAND PHARMA LTD	<u>40MG/2ML (20MG/ML)</u>	<u>A212993 001</u>	Nov 18, 2019
<u>AP</u>		<u>100MG/5ML (20MG/ML)</u>	<u>A212993 002</u>	Nov 18, 2019
<u>AP</u>	HENGRUI PHARMA	<u>40MG/2ML (20MG/ML)</u>	<u>A090675 002</u>	Dec 16, 2011
<u>AP</u>		<u>100MG/5ML (20MG/ML)</u>	<u>A090675 001</u>	Dec 16, 2011
<u>AP</u>	HIKMA	<u>40MG/2ML (20MG/ML)</u>	<u>A078753 001</u>	Dec 24, 2008
<u>AP</u>		<u>100MG/5ML (20MG/ML)</u>	<u>A078753 002</u>	Dec 24, 2008
<u>AP</u>	HIKMA FARMACEUTICA	<u>40MG/2ML (20MG/ML)</u>	<u>A091032 001</u>	Dec 20, 2010
<u>AP</u>		<u>100MG/5ML (20MG/ML)</u>	<u>A091032 002</u>	Dec 20, 2010
<u>AP</u>	HOSPIRA	<u>40MG/2ML (20MG/ML)</u>	<u>A077915 001</u>	Feb 27, 2008
<u>AP</u>		<u>100MG/5ML (20MG/ML)</u>	<u>A077915 002</u>	Feb 27, 2008
<u>AP</u>	!	<u>500MG/25ML (20MG/ML)</u>	<u>A078796 001</u>	Feb 27, 2008
<u>AP</u>	INTAS PHARMS USA	<u>40MG/2ML (20MG/ML)</u>	<u>A203054 001</u>	Aug 31, 2017
<u>AP</u>		<u>100MG/5ML (20MG/ML)</u>	<u>A203054 002</u>	Aug 31, 2017
<u>AP</u>	NOVAST LABS	<u>40MG/2ML (20MG/ML)</u>	<u>A206935 001</u>	May 26, 2017
<u>AP</u>		<u>100MG/5ML (20MG/ML)</u>	<u>A206935 002</u>	May 26, 2017
<u>AP</u>	QILU PHARM HAINAN	<u>40MG/2ML (20MG/ML)</u>	<u>A203380 001</u>	May 03, 2016
<u>AP</u>		<u>100MG/5ML (20MG/ML)</u>	<u>A203380 002</u>	May 03, 2016
<u>AP</u>		<u>300MG/15ML (20MG/ML)</u>	<u>A203380 003</u>	May 03, 2016
<u>AP</u>	SHILPA	<u>40MG/2ML (20MG/ML)</u>	<u>A208718 001</u>	Dec 28, 2018
<u>AP</u>		<u>100MG/5ML (20MG/ML)</u>	<u>A208718 002</u>	Dec 28, 2018
<u>AP</u>		<u>300MG/15ML (20MG/ML)</u>	<u>A208718 003</u>	Aug 16, 2019
<u>AP</u>	TEVA PHARMS USA	<u>500MG/25ML (20MG/ML)</u>	<u>A090101 001</u>	Nov 26, 2008
<u>AP</u>	ZENNOVA	<u>40MG/2ML (20MG/ML)</u>	<u>A090393 002</u>	May 13, 2011
<u>AP</u>		<u>100MG/5ML (20MG/ML)</u>	<u>A090393 003</u>	May 13, 2011

INJECTABLE, LIPOSOMAL; INTRAVENOUS

ONIVYDE

+! IPSEN

EQ 43MG BASE/10ML (EQ 4.3MG BASE/ML)

N207793 001 Oct 22, 2015

ISAVUCONAZONIUM SULFATE

CAPSULE; ORAL

CRESEMBA

+ ASTELLAS

74.5MG

N207500 002 Nov 22, 2022

+!

186MG

N207500 001 Mar 06, 2015

POWDER; INTRAVENOUS

CRESEMBA

+! ASTELLAS

372MG

N207501 001 Mar 06, 2015

ISOCARBOXAZID

TABLET; ORAL

MARPLAN

+! VALIDUS PHARMS INC

10MG

N011961 001

ISOFLURANE

LIQUID; INHALATION

FORANEAN +! BAXTER HLTHCARE99.9%N017624 001ISOFLURANEAN HALOCARBON PRODS99.9%A075225 001 Oct 20, 1999AN PIRAMAL CRITICAL99.9%A074416 001 Sep 30, 1994AN PIRAMAL PHARMA99.9%A074502 001 Jun 27, 1995ISONIAZID

INJECTABLE; INJECTION

ISONIAZID

! SANDOZ

100MG/ML

A040648 001 Jul 05, 2005

SYRUP; ORAL

ISONIAZID

+! CMP PHARMA INC

50MG/5ML

A088235 001 Nov 10, 1983

PRESCRIPTION DRUG PRODUCT LIST

ISONIAZID

TABLET; ORAL

ISONIAZID

<u>AA</u>	!	GENUS	<u>100MG</u>	<u>A080936</u>	<u>001</u>	
<u>AA</u>	!		<u>300MG</u>	<u>A080937</u>	<u>002</u>	
<u>AA</u>		OMNIVIUM PHARMS	<u>100MG</u>	<u>A040090</u>	<u>001</u>	Jun 26, 1997
<u>AA</u>			<u>300MG</u>	<u>A040090</u>	<u>002</u>	Jun 26, 1997
<u>AA</u>		THEPHARMANETWORK LLC	<u>100MG</u>	<u>A202610</u>	<u>001</u>	Oct 29, 2014
<u>AA</u>			<u>300MG</u>	<u>A202610</u>	<u>002</u>	Oct 29, 2014

ISOPROTERENOL HYDROCHLORIDE

INJECTABLE; INJECTION

ISOPROTERENOL HYDROCHLORIDE

<u>AP</u>		AMNEAL	<u>0.2MG/ML</u>	<u>A210576</u>	<u>001</u>	Oct 17, 2018
<u>AP</u>		AMPHASTAR PHARMS INC	<u>0.2MG/ML</u>	<u>A210106</u>	<u>001</u>	Jun 18, 2018
<u>AP</u>		AMRING PHARMS	<u>0.2MG/ML</u>	<u>A211237</u>	<u>001</u>	May 19, 2021
<u>AP</u>		AVET LIFESCIENCES	<u>0.2MG/ML</u>	<u>A212189</u>	<u>001</u>	Nov 19, 2021
<u>AP</u>		GLAND PHARMA LTD	<u>0.2MG/ML</u>	<u>A217648</u>	<u>001</u>	Oct 03, 2024
<u>AP</u>		HIKMA	<u>0.2MG/ML</u>	<u>A211703</u>	<u>001</u>	Apr 26, 2023
<u>AP</u>		MICRO LABS	<u>0.2MG/ML</u>	<u>A210845</u>	<u>001</u>	Feb 16, 2021
<u>AP</u>	!	NEXUS	<u>0.2MG/ML</u>	<u>A206961</u>	<u>001</u>	Aug 02, 2017
<u>AP</u>		PENN LIFE	<u>0.2MG/ML</u>	<u>A215542</u>	<u>001</u>	Sep 20, 2022
<u>AP</u>		SOMERSET THERAPS LLC	<u>0.2MG/ML</u>	<u>A211590</u>	<u>001</u>	Aug 06, 2024

ISOSORBIDE DINITRATE

TABLET; ORAL

ISORDIL

<u>AB</u>	+	BAUSCH	<u>5MG</u>	<u>N012093</u>	<u>007</u>	Jul 29, 1988
<u>AB</u>	+	!	<u>40MG</u>	<u>N012093</u>	<u>001</u>	Jul 29, 1988

ISOSORBIDE DINITRATE

<u>AB</u>		ENDO OPERATIONS	<u>5MG</u>	<u>A086923</u>	<u>001</u>	Mar 12, 1987
<u>AB</u>			<u>10MG</u>	<u>A086925</u>	<u>001</u>	Mar 12, 1987
<u>AB</u>			<u>20MG</u>	<u>A087537</u>	<u>001</u>	Oct 02, 1987
<u>AB</u>	!		<u>30MG</u>	<u>A087946</u>	<u>001</u>	Jan 12, 1988
<u>AB</u>			<u>40MG</u>	<u>A211290</u>	<u>001</u>	Nov 23, 2021
<u>AB</u>		HIKMA INTL PHARMS	<u>5MG</u>	<u>A086067</u>	<u>001</u>	Oct 29, 1987
<u>AB</u>			<u>10MG</u>	<u>A086066</u>	<u>001</u>	Oct 29, 1987
<u>AB</u>			<u>20MG</u>	<u>A088088</u>	<u>001</u>	Nov 02, 1987
<u>AB</u>		RUBICON	<u>5MG</u>	<u>A215723</u>	<u>001</u>	Jul 08, 2022
<u>AB</u>			<u>10MG</u>	<u>A215723</u>	<u>002</u>	Jul 08, 2022
<u>AB</u>			<u>20MG</u>	<u>A215723</u>	<u>003</u>	Jul 08, 2022
<u>AB</u>			<u>30MG</u>	<u>A215723</u>	<u>004</u>	Jul 08, 2022
<u>AB</u>			<u>40MG</u>	<u>A215723</u>	<u>005</u>	Jul 08, 2022
<u>AB</u>		SANDOZ	<u>5MG</u>	<u>A086221</u>	<u>001</u>	Jan 07, 1988
<u>AB</u>			<u>10MG</u>	<u>A086223</u>	<u>001</u>	Jan 07, 1988
<u>AB</u>			<u>20MG</u>	<u>A089367</u>	<u>001</u>	Apr 07, 1988
<u>AB</u>		ZYDUS LIFESCIENCES	<u>5MG</u>	<u>A213057</u>	<u>001</u>	Nov 20, 2019
<u>AB</u>			<u>10MG</u>	<u>A213057</u>	<u>002</u>	Nov 20, 2019
<u>AB</u>			<u>20MG</u>	<u>A213057</u>	<u>003</u>	Nov 20, 2019
<u>AB</u>			<u>30MG</u>	<u>A213057</u>	<u>004</u>	Nov 20, 2019
<u>AB</u>			<u>40MG</u>	<u>A213057</u>	<u>005</u>	Nov 20, 2019

ISOSORBIDE MONONITRATE

TABLET; ORAL

ISOSORBIDE MONONITRATE

<u>AB</u>		GENUS	<u>10MG</u>	<u>A075037</u>	<u>002</u>	Oct 30, 1998
<u>AB</u>			<u>20MG</u>	<u>A075037</u>	<u>001</u>	Oct 30, 1998

MONOKET

<u>AB</u>	+	OMNIVIUM PHARMS	<u>10MG</u>	<u>N020215</u>	<u>002</u>	Jun 30, 1993
<u>AB</u>	+	!	<u>20MG</u>	<u>N020215</u>	<u>001</u>	Jun 30, 1993

TABLET, EXTENDED RELEASE; ORAL

ISOSORBIDE MONONITRATE

<u>AB</u>		AUROBINDO PHARMA	<u>30MG</u>	<u>A216557</u>	<u>001</u>	Nov 07, 2022
<u>AB</u>			<u>60MG</u>	<u>A216557</u>	<u>002</u>	Nov 07, 2022
<u>AB</u>			<u>120MG</u>	<u>A216557</u>	<u>003</u>	Nov 07, 2022
<u>AB</u>		CHARTWELL MOLECULAR	<u>30MG</u>	<u>A075155</u>	<u>002</u>	Jan 13, 2000
<u>AB</u>			<u>60MG</u>	<u>A075155</u>	<u>001</u>	Oct 30, 1998
<u>AB</u>			<u>120MG</u>	<u>A075155</u>	<u>003</u>	Aug 04, 2000
<u>AB</u>		DEXCEL LTD	<u>30MG</u>	<u>A075522</u>	<u>002</u>	Sep 20, 2016
<u>AB</u>			<u>60MG</u>	<u>A075522</u>	<u>001</u>	Apr 17, 2000
<u>AB</u>			<u>120MG</u>	<u>A210822</u>	<u>001</u>	Aug 29, 2018

PRESCRIPTION DRUG PRODUCT LIST

ISOSORBIDE MONONITRATE

TABLET, EXTENDED RELEASE;ORAL

ISOSORBIDE MONONITRATE

<u>AB</u>	RICONPHARMA LLC	<u>30MG</u>	<u>A210918 001</u>	Nov 05, 2018
<u>AB</u>		<u>60MG</u>	<u>A210918 002</u>	Nov 05, 2018
<u>AB</u>	!	<u>120MG</u>	<u>A210918 003</u>	Nov 05, 2018
<u>AB</u>	SHANDONG	<u>30MG</u>	<u>A214115 002</u>	Feb 08, 2024
<u>AB</u>		<u>60MG</u>	<u>A214115 003</u>	Feb 08, 2024
<u>AB</u>		<u>120MG</u>	<u>A214115 001</u>	Apr 29, 2022
<u>AB</u>	TORRENT PHARMS	<u>30MG</u>	<u>A200270 001</u>	Jun 03, 2011
<u>AB</u>		<u>60MG</u>	<u>A200495 001</u>	Jun 03, 2011
<u>AB</u>		<u>120MG</u>	<u>A200495 002</u>	Jun 03, 2011
<u>AB</u>	ZYDUS HLTHCARE	<u>30MG</u>	<u>A075395 001</u>	Mar 16, 2000
<u>AB</u>		<u>60MG</u>	<u>A075395 002</u>	Mar 16, 2000
<u>AB</u>		<u>120MG</u>	<u>A075395 003</u>	Mar 16, 2000

ISOSULFAN BLUE

SOLUTION;SUBCUTANEOUS

ISOSULFAN BLUE

<u>AP</u>	AM REGENT	<u>50MG/5ML (10MG/ML)</u>	<u>A210294 001</u>	May 17, 2023
<u>AP</u>	MEITHEAL	<u>50MG/5ML (10MG/ML)</u>	<u>A213130 001</u>	Nov 03, 2021
<u>AP</u>	MSN	<u>50MG/5ML (10MG/ML)</u>	<u>A216090 001</u>	Sep 16, 2022
<u>AP</u>	! MYLAN INSTITUTIONAL	<u>50MG/5ML (10MG/ML)</u>	<u>A090874 001</u>	Jul 20, 2010

ISOTRETINOIN

CAPSULE;ORAL

AMNESTEEM

<u>AB1</u>	MYLAN PHARMS INC	<u>10MG</u>	<u>A075945 001</u>	Nov 08, 2002
<u>AB1</u>		<u>20MG</u>	<u>A075945 002</u>	Nov 08, 2002
<u>AB1</u>		<u>40MG</u>	<u>A075945 003</u>	Nov 08, 2002

CLARAVIS

<u>AB1</u>	TEVA PHARMS USA	<u>10MG</u>	<u>A076356 001</u>	Apr 11, 2003
<u>AB1</u>		<u>20MG</u>	<u>A076135 002</u>	Apr 11, 2003
<u>AB1</u>		<u>30MG</u>	<u>A076135 003</u>	May 11, 2006
<u>AB1</u>	!	<u>40MG</u>	<u>A076135 001</u>	Apr 11, 2003

ISOTRETINOIN

<u>AB1</u>	AMNEAL PHARMS NY	<u>10MG</u>	<u>A207792 001</u>	Sep 29, 2017
<u>AB1</u>		<u>20MG</u>	<u>A207792 002</u>	Sep 29, 2017
<u>AB1</u>		<u>30MG</u>	<u>A207792 003</u>	Sep 29, 2017
<u>AB1</u>		<u>40MG</u>	<u>A207792 004</u>	Sep 29, 2017
<u>AB1</u>	ZYDUS PHARMS	<u>10MG</u>	<u>A211568 001</u>	Aug 29, 2023
<u>AB1</u>		<u>20MG</u>	<u>A211568 002</u>	Aug 29, 2023
<u>AB1</u>		<u>30MG</u>	<u>A211568 003</u>	Aug 29, 2023
<u>AB1</u>		<u>40MG</u>	<u>A211568 004</u>	Aug 29, 2023

MYORISAN

<u>AB1</u>	UPSHER SMITH LABS	<u>10MG</u>	<u>A076485 001</u>	Jan 19, 2012
<u>AB1</u>		<u>20MG</u>	<u>A076485 002</u>	Jan 19, 2012
<u>AB1</u>		<u>30MG</u>	<u>A076485 004</u>	Aug 25, 2015
<u>AB1</u>		<u>40MG</u>	<u>A076485 003</u>	Jan 19, 2012

ZENATANE

<u>AB1</u>	DR REDDYS LABS LTD	<u>10MG</u>	<u>A202099 001</u>	Mar 25, 2013
<u>AB1</u>		<u>20MG</u>	<u>A202099 002</u>	Mar 25, 2013
<u>AB1</u>		<u>30MG</u>	<u>A202099 004</u>	Feb 23, 2015
<u>AB1</u>		<u>40MG</u>	<u>A202099 003</u>	Mar 25, 2013

ABSORICA

<u>AB2</u>	+ SUN PHARM INDS INC	<u>10MG</u>	<u>N021951 001</u>	May 25, 2012
<u>AB2</u>	+	<u>20MG</u>	<u>N021951 002</u>	May 25, 2012
<u>AB2</u>	+	<u>25MG</u>	<u>N021951 005</u>	Aug 15, 2014
<u>AB2</u>	+	<u>30MG</u>	<u>N021951 003</u>	May 25, 2012
<u>AB2</u>	+	<u>35MG</u>	<u>N021951 006</u>	Aug 15, 2014
<u>AB2</u>	+!	<u>40MG</u>	<u>N021951 004</u>	May 25, 2012

ISOTRETINOIN

<u>AB2</u>	ACTAVIS LABS FL	<u>10MG</u>	<u>A205063 001</u>	Mar 31, 2021
<u>AB2</u>		<u>20MG</u>	<u>A205063 002</u>	Mar 31, 2021
<u>AB2</u>		<u>25MG</u>	<u>A205063 003</u>	Mar 31, 2021
<u>AB2</u>		<u>30MG</u>	<u>A205063 004</u>	Mar 31, 2021
<u>AB2</u>		<u>35MG</u>	<u>A205063 005</u>	Mar 31, 2021
<u>AB2</u>		<u>40MG</u>	<u>A205063 006</u>	Mar 31, 2021
<u>AB2</u>	AUROBINDO PHARMA	<u>10MG</u>	<u>A218194 001</u>	Jan 29, 2024
<u>AB2</u>		<u>20MG</u>	<u>A218194 002</u>	Jan 29, 2024
<u>AB2</u>		<u>25MG</u>	<u>A218194 003</u>	Jan 29, 2024
<u>AB2</u>		<u>30MG</u>	<u>A218194 004</u>	Jan 29, 2024
<u>AB2</u>		<u>35MG</u>	<u>A218194 005</u>	Jan 29, 2024

PRESCRIPTION DRUG PRODUCT LIST

ISOTRETINOIN

CAPSULE; ORAL

ISOTRETINOIN

<u>AB2</u>		<u>40MG</u>	<u>A218194</u>	<u>006</u>	Jan 29, 2024
<u>AB2</u>	UPSHER SMITH LABS	<u>10MG</u>	<u>A212333</u>	<u>001</u>	Sep 21, 2021
<u>AB2</u>		<u>20MG</u>	<u>A212333</u>	<u>002</u>	Sep 21, 2021
<u>AB2</u>		<u>30MG</u>	<u>A212333</u>	<u>003</u>	Sep 21, 2021
<u>AB2</u>		<u>40MG</u>	<u>A213571</u>	<u>001</u>	Apr 12, 2021
	ABSORICA LD				
	+ SUN PHARM	8MG	N211913	001	Nov 05, 2019
	+	16MG	N211913	002	Nov 05, 2019
	+	24MG	N211913	004	Nov 05, 2019
	+	32MG	N211913	006	Nov 05, 2019

ISRADIPINE

CAPSULE; ORAL

ISRADIPINE

<u>AB</u>	ELITE LABS INC	<u>2.5MG</u>	<u>A077169</u>	<u>001</u>	Apr 24, 2006
<u>AB</u>		<u>5MG</u>	<u>A077169</u>	<u>002</u>	Apr 24, 2006
<u>AB</u>	WATSON LABS TEVA	<u>2.5MG</u>	<u>A077317</u>	<u>001</u>	Jan 05, 2006
<u>AB</u>	!	<u>5MG</u>	<u>A077317</u>	<u>002</u>	Jan 05, 2006

ISTRADIFYLLINE

TABLET; ORAL

ISTRADIFYLLINE

	NOURIANZ				
	+ KYOWA KIRIN	20MG	N022075	001	Aug 27, 2019
	+	40MG	N022075	002	Aug 27, 2019

ITRACONAZOLE

CAPSULE; ORAL

ITRACONAZOLE

<u>AB</u>	ALEMBIC	<u>100MG</u>	<u>A206741</u>	<u>001</u>	Dec 13, 2016
<u>AB</u>	ALKEM LABS LTD	<u>100MG</u>	<u>A208591</u>	<u>001</u>	Jun 12, 2017
<u>AB</u>	AMNEAL PHARMS	<u>100MG</u>	<u>A205080</u>	<u>001</u>	Sep 26, 2016
<u>AB</u>	SANDOZ	<u>100MG</u>	<u>A076104</u>	<u>001</u>	May 28, 2004
<u>AB</u>	TORRENT	<u>100MG</u>	<u>A209460</u>	<u>001</u>	Aug 24, 2018
<u>AB</u>	ZYDUS PHARMS	<u>100MG</u>	<u>A204672</u>	<u>001</u>	Sep 19, 2017

SPORANOX

<u>AB</u>	+	JANSSEN PHARMS	<u>100MG</u>	<u>N020083</u>	<u>001</u>	Sep 11, 1992
	TOLSURA					
	+	MAYNE PHARMA	65MG	N208901	001	Dec 11, 2018

SOLUTION; ORAL

ITRACONAZOLE

<u>AA</u>	AMNEAL PHARMS	<u>10MG/ML</u>	<u>A205573</u>	<u>001</u>	Oct 30, 2015
<u>AA</u>	ANNORA PHARMA	<u>10MG/ML</u>	<u>A212239</u>	<u>001</u>	Sep 01, 2020

SPORANOX

<u>AA</u>	+	JANSSEN PHARMS	<u>10MG/ML</u>	<u>N020657</u>	<u>001</u>	Feb 21, 1997
-----------	---	----------------	----------------	----------------	------------	--------------

IVABRADINE

SOLUTION; ORAL

IVABRADINE

	CORLANOR					
	+	AMGEN INC	5MG/5ML (1MG/ML)	N209964	001	Apr 22, 2019

IVABRADINE HYDROCHLORIDE

TABLET; ORAL

CORLANOR

<u>AB</u>	+	AMGEN INC	<u>EQ 5MG BASE</u>	<u>N206143</u>	<u>001</u>	Apr 15, 2015
<u>AB</u>	+		<u>EQ 7.5MG BASE</u>	<u>N206143</u>	<u>002</u>	Apr 15, 2015

IVABRADINE HYDROCHLORIDE

<u>AB</u>	ALEMBIC	<u>EQ 5MG BASE</u>	<u>A215238</u>	<u>001</u>	Nov 08, 2024
<u>AB</u>		<u>EQ 7.5MG BASE</u>	<u>A215238</u>	<u>002</u>	Nov 08, 2024
<u>AB</u>	ANNORA PHARMA	<u>EQ 5MG BASE</u>	<u>A213366</u>	<u>001</u>	Oct 05, 2022
<u>AB</u>		<u>EQ 7.5MG BASE</u>	<u>A213366</u>	<u>002</u>	Oct 05, 2022
<u>AB</u>	BIONPHARMA	<u>EQ 5MG BASE</u>	<u>A213276</u>	<u>001</u>	Sep 05, 2024
<u>AB</u>		<u>EQ 7.5MG BASE</u>	<u>A213276</u>	<u>002</u>	Sep 05, 2024
<u>AB</u>	INGENUS PHARMS LLC	<u>EQ 5MG BASE</u>	<u>A214051</u>	<u>001</u>	Dec 30, 2021
<u>AB</u>		<u>EQ 7.5MG BASE</u>	<u>A214051</u>	<u>002</u>	Dec 30, 2021
<u>AB</u>	ZYDUS PHARMS	<u>EQ 5MG BASE</u>	<u>A213442</u>	<u>001</u>	Nov 29, 2023
<u>AB</u>		<u>EQ 7.5MG BASE</u>	<u>A213442</u>	<u>002</u>	Nov 29, 2023

PRESCRIPTION DRUG PRODUCT LIST

IVACAFTOR

GRANULE; ORAL

KALYDECO

+	VERTEX PHARMS INC	5.8MG/PACKET	N207925	004	May 03, 2023
+		13.4MG/PACKET	N207925	005	May 03, 2023
+		25MG/PACKET	N207925	003	Apr 29, 2019
+		50MG/PACKET	N207925	001	Mar 17, 2015
+	!	75MG/PACKET	N207925	002	Mar 17, 2015

TABLET; ORAL

KALYDECO

+	!	VERTEX PHARMS	150MG	N203188	001	Jan 31, 2012
---	---	---------------	-------	---------	-----	--------------

IVACAFTOR; IVACAFTOR, TEZACAFTOR

TABLET; ORAL

SYMDEKO (COPACKAGED)

+	VERTEX PHARMS INC	75MG;75MG, 50MG	N210491	002	Jun 21, 2019
+	!	150MG;150MG, 100MG	N210491	001	Feb 12, 2018

IVACAFTOR; LUMACAFTOR

GRANULE; ORAL

ORKAMBI

+	VERTEX PHARMS INC	94MG/PACKET;75MG/PACKET	N211358	003	Sep 02, 2022
+		125MG/PACKET;100MG/PACKET	N211358	001	Aug 07, 2018
+	!	188MG/PACKET;150MG/PACKET	N211358	002	Aug 07, 2018

TABLET; ORAL

ORKAMBI

+	VERTEX PHARMS INC	125MG;100MG	N206038	002	Sep 28, 2016
+	!	125MG;200MG	N206038	001	Jul 02, 2015

IVERMECTIN

CREAM; TOPICAL

IVERMECTIN

AB	+	PADAGIS ISRAEL	1%	A210225	001	Apr 13, 2020
AB	+	TEVA PHARMS USA	1%	A210019	001	Sep 13, 2019
AB	+	ZYDUS LIFESCIENCES	1%	A215210	001	Aug 02, 2022

SOOLANTRA

AB	+	!	GALDERMA LABS LP	1%	N206255	001	Dec 19, 2014
-----------	---	---	------------------	-----------	----------------	------------	--------------

TABLET; ORAL

IVERMECTIN

AB		EDENBRIDGE PHARMS	3MG	A204154	001	Oct 24, 2014
AB		SENORES PHARMS	3MG	A218324	001	Oct 16, 2024

STROMECTOL

AB	+	!	MERCK SHARP DOHME	3MG	N050742	002	Oct 08, 1998
-----------	---	---	-------------------	------------	----------------	------------	--------------

IVOSIDENIB

TABLET; ORAL

TIBSOVO

+	!	SERVIER	250MG	N211192	001	Jul 20, 2018
---	---	---------	-------	---------	-----	--------------

IXABEPILONE

INJECTABLE; INTRAVENOUS

IXEMPRA KIT

+	!	R-PHARM US LLC	15MG/VIAL	N022065	001	Oct 16, 2007
+	!		45MG/VIAL	N022065	002	Oct 16, 2007

IXAZOMIB CITRATE

CAPSULE; ORAL

NINLARO

+	TAKEDA PHARMS USA	EQ 2.3MG BASE	N208462	001	Nov 20, 2015
+		EQ 3MG BASE	N208462	002	Nov 20, 2015
+	!	EQ 4MG BASE	N208462	003	Nov 20, 2015

KETAMINE HYDROCHLORIDE

INJECTABLE; INJECTION

KETALAR

AP	+	!	ENDO OPERATIONS	EQ 10MG BASE/ML	N016812	001
AP	+			EQ 50MG BASE/ML	N016812	002
AP	+			EQ 100MG BASE/ML	N016812	003

KETAMINE HYDROCHLORIDE

AP		EUGIA PHARMA	EQ 10MG BASE/ML	A076092	001	Sep 30, 2008
AP			EQ 50MG BASE/ML	A076092	002	Dec 28, 2001
AP			EQ 100MG BASE/ML	A076092	003	Oct 25, 2002
AP		FRESENIUS KABI USA	EQ 10MG BASE/ML	A215808	001	Jan 13, 2023
AP			EQ 50MG BASE/ML	A215808	002	Jan 13, 2023
AP		GLAND PHARMA LTD	EQ 10MG BASE/ML	A216809	001	Jan 24, 2023
AP			EQ 50MG BASE/ML	A216809	002	Jan 24, 2023

PRESCRIPTION DRUG PRODUCT LIST

KETAMINE HYDROCHLORIDE

INJECTABLE; INJECTION

KETAMINE HYDROCHLORIDE

<u>AP</u>		<u>EQ 100MG BASE/ML</u>	<u>A216809 003</u>	Jan 24, 2023	
<u>AP</u>	!	HIKMA	<u>EQ 50MG BASE/ML</u>	<u>A074524 001</u>	Mar 22, 1996
<u>AP</u>	!		<u>EQ 100MG BASE/ML</u>	<u>A074524 002</u>	Mar 22, 1996
<u>AP</u>		HOSPIRA	<u>EQ 50MG BASE/ML</u>	<u>A074549 001</u>	Jun 27, 1996
<u>AP</u>			<u>EQ 100MG BASE/ML</u>	<u>A074549 002</u>	Jun 27, 1996

KETOCONAZOLE

AEROSOL, FOAM; TOPICAL

KETOCONAZOLE

<u>AB</u>		PADAGIS ISRAEL	<u>2%</u>	<u>A091550 001</u>	Aug 25, 2011
<u>AB</u>	!	XIROMED	<u>2%</u>	<u>A213601 001</u>	May 21, 2021

CREAM; TOPICAL

KETOCONAZOLE

<u>AB</u>		ENCUBE	<u>2%</u>	<u>A212443 001</u>	May 20, 2021
<u>AB</u>		FOUGERA PHARMS	<u>2%</u>	<u>A076294 001</u>	Apr 28, 2004
<u>AB</u>	!	TEVA	<u>2%</u>	<u>A075581 001</u>	Apr 25, 2000
<u>AB</u>		TRUPHARMA	<u>2%</u>	<u>A215185 001</u>	Nov 17, 2021

KETOZOLE

<u>AB</u>		TARO	<u>2%</u>	<u>A075638 001</u>	Dec 18, 2002
-----------	--	------	-----------	--------------------	--------------

SHAMPOO; TOPICAL

KETOCONAZOLE

<u>AB</u>		COSETTE	<u>2%</u>	<u>A076942 001</u>	Apr 11, 2005
<u>AB</u>		NOVITIUM PHARMA	<u>2%</u>	<u>A218498 001</u>	Sep 16, 2024
<u>AB</u>	!	PADAGIS ISRAEL	<u>2%</u>	<u>A076419 001</u>	Jan 07, 2004

TABLET; ORAL

KETOCONAZOLE

<u>AB</u>		SENORES PHARMS	<u>200MG</u>	<u>A075912 001</u>	Jan 10, 2002
<u>AB</u>		STRIDES PHARMA	<u>200MG</u>	<u>A210457 001</u>	Jun 18, 2018
<u>AB</u>	!	TARO	<u>200MG</u>	<u>A075319 001</u>	Jun 15, 1999

KETOPROFEN

CAPSULE; ORAL

KETOPROFEN

<u>AB</u>		MISEMER	<u>50MG</u>	<u>A074014 002</u>	Jan 29, 1993
<u>AB</u>		TEVA	<u>50MG</u>	<u>A073516 001</u>	Dec 22, 1992
		MISEMER	25MG	A074014 001	Jan 29, 1993
	!		75MG	A074014 003	Jan 29, 1993

CAPSULE, EXTENDED RELEASE; ORAL

KETOPROFEN

!	MYLAN	200MG	A075679 001	Feb 20, 2002
---	-------	-------	-------------	--------------

KETOROLAC TROMETHAMINE

INJECTABLE; INJECTION

KETOROLAC TROMETHAMINE

<u>AP</u>		ALEMBIC	<u>15MG/ML</u>	<u>A214456 001</u>	Nov 02, 2022
<u>AP</u>			<u>30MG/ML</u>	<u>A214456 002</u>	Nov 02, 2022
<u>AP</u>		ASPIRO	<u>15MG/ML</u>	<u>A217166 001</u>	Jun 20, 2023
<u>AP</u>			<u>30MG/ML</u>	<u>A217166 002</u>	Jun 20, 2023
<u>AP</u>		BAXTER HLTHCARE CORP	<u>15MG/ML</u>	<u>A209900 002</u>	Jul 25, 2018
<u>AP</u>			<u>30MG/ML</u>	<u>A209900 001</u>	Sep 15, 2017
<u>AP</u>		CAPLIN	<u>15MG/ML</u>	<u>A217789 001</u>	May 10, 2023
<u>AP</u>			<u>30MG/ML</u>	<u>A217789 002</u>	May 10, 2023
<u>AP</u>		FRESENIUS KABI USA	<u>15MG/ML</u>	<u>A075784 001</u>	Jan 11, 2002
<u>AP</u>			<u>15MG/ML</u>	<u>A203242 001</u>	Oct 07, 2015
<u>AP</u>			<u>30MG/ML</u>	<u>A075784 002</u>	Jan 11, 2002
<u>AP</u>			<u>30MG/ML</u>	<u>A203242 002</u>	Oct 07, 2015
<u>AP</u>		GLAND	<u>15MG/ML</u>	<u>A204216 001</u>	Nov 01, 2016
<u>AP</u>			<u>30MG/ML</u>	<u>A204216 002</u>	Nov 01, 2016
<u>AP</u>		HIKMA	<u>15MG/ML</u>	<u>A075772 001</u>	Jul 21, 2004
<u>AP</u>			<u>30MG/ML</u>	<u>A075772 002</u>	Jul 21, 2004
<u>AP</u>	!	HOSPIRA	<u>15MG/ML</u>	<u>A074802 001</u>	Jun 05, 1997
<u>AP</u>	!		<u>30MG/ML</u>	<u>A074802 002</u>	Jun 05, 1997
<u>AP</u>			<u>30MG/ML</u>	<u>A074993 002</u>	Jan 27, 1999
<u>AP</u>		NEPHRON	<u>30MG/ML</u>	<u>A211445 001</u>	Aug 20, 2020
<u>AP</u>		SAGENT PHARMS INC	<u>15MG/ML</u>	<u>A091065 001</u>	Nov 27, 2013
<u>AP</u>			<u>30MG/ML</u>	<u>A091065 002</u>	Nov 27, 2013
<u>AP</u>		SANDOZ	<u>30MG/ML</u>	<u>A076271 002</u>	Oct 06, 2004
<u>AP</u>		SUN PHARM	<u>15MG/ML</u>	<u>A078737 001</u>	Oct 06, 2008
<u>AP</u>			<u>30MG/ML</u>	<u>A078737 002</u>	Oct 06, 2008
<u>AP</u>		WOCKHARDT	<u>15MG/ML</u>	<u>A077942 001</u>	Mar 27, 2007

PRESCRIPTION DRUG PRODUCT LIST

KETOROLAC TROMETHAMINE

INJECTABLE; INJECTION

KETOROLAC TROMETHAMINE

AP **30MG/ML** **A077942 002** Mar 27, 2007
SOLUTION/DROPS;OPHTHALMIC

ACULAR

AT **+!** **ABBVIE** **0.5%** **N019700 001** Nov 09, 1992

ACULAR LS

AT **+!** **ABBVIE** **0.4%** **N021528 001** May 30, 2003

KETOROLAC TROMETHAMINE

AT **APOTEX INC** **0.4%** **A077308 001** Nov 05, 2009

AT **0.5%** **A076109 001** Nov 05, 2009

AT **CAPLIN** **0.5%** **A218204 001** Mar 26, 2024

AT **MICRO LABS LTD** **0.5%** **A203410 001** Apr 05, 2019

INDIA

AT **SANDOZ** **0.5%** **A076583 001** Nov 05, 2009

ACUVAIL

+! **ABBVIE** **0.45%** **N022427 001** Jul 22, 2009

SPRAY, METERED;NASAL

SPRIX

+! **ZYLA** **15.75MG/SPRAY** **N022382 001** May 14, 2010

TABLET;ORAL

KETOROLAC TROMETHAMINE

AB **ATNAHS PHARMA US** **10MG** **A216407 001** Oct 21, 2022

AB **BIONPHARMA** **10MG** **A216759 001** Feb 23, 2023

AB **CHARTWELL RX** **10MG** **A210616 001** Aug 16, 2018

AB **HETERO LABS LTD III** **10MG** **A216651 001** Aug 25, 2022

AB **LEADING** **10MG** **A215745 001** Feb 23, 2023

AB **MYLAN** **10MG** **A074761 001** May 16, 1997

AB **SENORES PHARMS** **10MG** **A215788 001** Mar 23, 2022

AB **!** **TEVA** **10MG** **A074754 001** May 16, 1997

AB **ZYDUS LIFESCIENCES** **10MG** **A217038 001** Oct 21, 2022

KETOROLAC TROMETHAMINE; PHENYLEPHRINE HYDROCHLORIDE

SOLUTION;IRRIGATION

KETOROLAC TROMETHAMINE AND PHENYLEPHRINE HYDROCHLORIDE

AT **LUPIN LTD** **EQ 0.3% BASE;EQ 1% BASE** **A210183 001** Jul 01, 2019

OMIDRIA

AT **+!** **RAYNER SURGICAL** **EQ 0.3% BASE;EQ 1% BASE** **N205388 001** May 30, 2014

L-GLUTAMINE

FOR SOLUTION;ORAL

ENDARI

AA **+** **EMMAUS MEDCL** **5GM/PACKET** **N208587 001** Jul 07, 2017

L-GLUTAMINE

AA **NOVITIUM PHARMA** **5GM/PACKET** **A215647 001** Jul 08, 2024

LABELTALOL HYDROCHLORIDE

INJECTABLE; INJECTION

LABELTALOL HYDROCHLORIDE

AP **BAXTER HLTHCARE** **5MG/ML** **A076051 001** Jul 05, 2002

CORP

AP **CAPLIN** **5MG/ML** **A214533 001** Sep 07, 2021

AP **GLAND PHARMA LTD** **5MG/ML** **A090699 001** Apr 03, 2012

AP **HIKMA** **5MG/ML** **A075303 001** May 28, 1999

AP **!** **HOSPIRA** **5MG/ML** **A075239 001** Nov 29, 1999

AP **!** **5MG/ML** **A075240 001** Nov 29, 1999

AP **RISING** **5MG/ML** **A075431 001** Nov 29, 1999

SOLUTION;INTRAVENOUS

LABELTALOL HYDROCHLORIDE

+! **HIKMA** **20MG/4ML (5MG/ML)** **N213330 006** Aug 19, 2024

LABELTALOL HYDROCHLORIDE IN DEXTROSE

+! **HIKMA** **200MG/200ML (1MG/ML)** **N213330 001** Nov 09, 2020

LABELTALOL HYDROCHLORIDE IN SODIUM CHLORIDE

+! **HIKMA** **100MG/100ML (1MG/ML)** **N213330 002** Nov 09, 2020

+! **200MG/200ML (1MG/ML)** **N213330 003** Nov 09, 2020

+! **300MG/300ML (1MG/ML)** **N213330 004** Nov 09, 2020

TABLET;ORAL

LABELTALOL HYDROCHLORIDE

AB **ANDAS 5 HOLDING** **100MG** **A075215 001** Jul 29, 1999

AB **200MG** **A075215 002** Jul 29, 1999

AB **300MG** **A075215 003** Jul 29, 1999

AB **APPCO** **100MG** **A209603 001** Jun 20, 2018

AB **200MG** **A209603 002** Jun 20, 2018

PRESCRIPTION DRUG PRODUCT LIST

LABETALOL HYDROCHLORIDE

TABLET; ORAL

LABETALOL HYDROCHLORIDE

<u>AB</u>		<u>300MG</u>	<u>A209603 003</u>	Jun 20, 2018
<u>AB</u>	CADILA PHARMS LTD	<u>100MG</u>	<u>A211325 001</u>	May 13, 2019
<u>AB</u>		<u>200MG</u>	<u>A211325 002</u>	May 13, 2019
<u>AB</u>		<u>300MG</u>	<u>A211325 003</u>	May 13, 2019
<u>AB</u>	ENDO OPERATIONS	<u>100MG</u>	<u>A200908 001</u>	Jul 10, 2012
<u>AB</u>	!	<u>200MG</u>	<u>A200908 002</u>	Jul 10, 2012
<u>AB</u>		<u>300MG</u>	<u>A200908 003</u>	Jul 10, 2012
<u>AB</u>	EPIC PHARMA LLC	<u>100MG</u>	<u>A212990 001</u>	Sep 30, 2020
<u>AB</u>		<u>200MG</u>	<u>A212990 002</u>	Sep 30, 2020
<u>AB</u>		<u>300MG</u>	<u>A212990 003</u>	Sep 30, 2020
<u>AB</u>	HERITAGE PHARMA	<u>100MG</u>	<u>A074787 001</u>	Aug 03, 1998
<u>AB</u>		<u>200MG</u>	<u>A074787 002</u>	Aug 03, 1998
<u>AB</u>		<u>300MG</u>	<u>A074787 003</u>	Aug 03, 1998
<u>AB</u>	HIBROW HLTHCARE	<u>100MG</u>	<u>A207863 001</u>	Feb 04, 2019
<u>AB</u>		<u>200MG</u>	<u>A207863 002</u>	Feb 04, 2019
<u>AB</u>		<u>300MG</u>	<u>A207863 003</u>	Feb 04, 2019
<u>AB</u>	RUBICON	<u>100MG</u>	<u>A211953 001</u>	Aug 18, 2021
<u>AB</u>		<u>200MG</u>	<u>A211953 002</u>	Aug 18, 2021
<u>AB</u>		<u>300MG</u>	<u>A211953 003</u>	Aug 18, 2021
<u>AB</u>	SANDOZ	<u>100MG</u>	<u>A075113 001</u>	Aug 04, 1998
<u>AB</u>	!	<u>200MG</u>	<u>A075113 002</u>	Aug 04, 1998
<u>AB</u>		<u>300MG</u>	<u>A075113 003</u>	Aug 04, 1998
<u>AB</u>	WATSON LABS	<u>100MG</u>	<u>A075133 001</u>	Aug 03, 1998
<u>AB</u>		<u>200MG</u>	<u>A075133 002</u>	Aug 03, 1998
<u>AB</u>		<u>300MG</u>	<u>A075133 003</u>	Aug 03, 1998
<u>AB</u>	ZYDUS PHARMS	<u>100MG</u>	<u>A207743 001</u>	Sep 19, 2017
<u>AB</u>		<u>200MG</u>	<u>A207743 002</u>	Sep 19, 2017
<u>AB</u>		<u>300MG</u>	<u>A207743 003</u>	Sep 19, 2017
	APPCO	400MG	A209603 004	Oct 18, 2024

LACOSAMIDE

CAPSULE, EXTENDED RELEASE; ORAL

MOTPOLY XR

+	AUCTA	100MG	N216185 001	May 04, 2023
+		150MG	N216185 002	May 04, 2023
+	!	200MG	N216185 003	May 04, 2023

SOLUTION; INTRAVENOUS

LACOSAMIDE

<u>AP</u>	ACELLA	<u>200MG/20ML (10MG/ML)</u>	<u>A218464 001</u>	Nov 14, 2024
<u>AP</u>	APOTEX	<u>200MG/20ML (10MG/ML)</u>	<u>A216670 001</u>	Nov 13, 2023
<u>AP</u>	ASPIRO	<u>200MG/20ML (10MG/ML)</u>	<u>A216335 001</u>	Sep 14, 2022
<u>AP</u>	DR REDDYS	<u>200MG/20ML (10MG/ML)</u>	<u>A217718 001</u>	Mar 25, 2024
<u>AP</u>	FRESENIUS KABI USA	<u>200MG/20ML (10MG/ML)</u>	<u>A214677 001</u>	May 03, 2022
<u>AP</u>	GLAND PHARMA LTD	<u>200MG/20ML (10MG/ML)</u>	<u>A215628 001</u>	Sep 19, 2022
<u>AP</u>	HAINAN POLY	<u>200MG/20ML (10MG/ML)</u>	<u>A217311 001</u>	Feb 13, 2024
<u>AP</u>	HIKMA	<u>200MG/20ML (10MG/ML)</u>	<u>A216806 001</u>	Apr 05, 2024
<u>AP</u>	INDOCO	<u>200MG/20ML (10MG/ML)</u>	<u>A214301 001</u>	Apr 07, 2022
<u>AP</u>	MSN	<u>200MG/20ML (10MG/ML)</u>	<u>A215979 001</u>	Feb 09, 2023
<u>AP</u>	SOMERSET THERAPS LLC	<u>200MG/20ML (10MG/ML)</u>	<u>A214960 001</u>	Oct 09, 2024
<u>AP</u>	ZYDUS PHARMS	<u>200MG/20ML (10MG/ML)</u>	<u>A209465 001</u>	Jun 29, 2022

VIMPAT

<u>AP</u>	+	!	UCB INC	<u>200MG/20ML (10MG/ML)</u>	<u>N022254 001</u>	Oct 28, 2008
-----------	---	---	---------	-----------------------------	--------------------	--------------

SOLUTION; ORAL

LACOSAMIDE

<u>AA</u>	ALKEM LABS LTD	<u>10MG/ML</u>	<u>A214672 001</u>	May 19, 2022
<u>AA</u>	APOTEX	<u>10MG/ML</u>	<u>A206355 001</u>	Sep 26, 2022
<u>AA</u>	AUROBINDO PHARMA LTD	<u>10MG/ML</u>	<u>A209224 001</u>	Jan 24, 2024
<u>AA</u>	CHARTWELL RX	<u>10MG/ML</u>	<u>A215154 001</u>	May 15, 2024
<u>AA</u>	HETERO LABS LTD III	<u>10MG/ML</u>	<u>A209301 001</u>	May 31, 2022
<u>AA</u>	MEDLEY PHARMS	<u>10MG/ML</u>	<u>A216461 001</u>	Feb 06, 2023
<u>AA</u>	MSN	<u>10MG/ML</u>	<u>A215379 001</u>	Dec 22, 2023
<u>AA</u>	NOVITIUM PHARMA	<u>10MG/ML</u>	<u>A216151 001</u>	Aug 26, 2022

VIMPAT

<u>AA</u>	+	!	UCB INC	<u>10MG/ML</u>	<u>N022255 001</u>	Apr 20, 2010
-----------	---	---	---------	----------------	--------------------	--------------

TABLET; ORAL

LACOSAMIDE

<u>AB</u>	ALEMBIC	<u>50MG</u>	<u>A204974 001</u>	Mar 17, 2022
<u>AB</u>		<u>100MG</u>	<u>A204974 002</u>	Mar 17, 2022

PRESCRIPTION DRUG PRODUCT LIST

LACOSAMIDE

TABLET; ORAL

LACOSAMIDE

<u>AB</u>		<u>150MG</u>	<u>A204974 003</u>	Mar 17, 2022
<u>AB</u>		<u>200MG</u>	<u>A204974 004</u>	Mar 17, 2022
<u>AB</u>	ALKEM LABS LTD	<u>50MG</u>	<u>A214695 001</u>	Mar 31, 2022
<u>AB</u>		<u>100MG</u>	<u>A214695 002</u>	Mar 31, 2022
<u>AB</u>		<u>150MG</u>	<u>A214695 003</u>	Mar 31, 2022
<u>AB</u>		<u>200MG</u>	<u>A214695 004</u>	Mar 31, 2022
<u>AB</u>	AUROBINDO PHARMA LTD	<u>50MG</u>	<u>A204994 001</u>	Jan 05, 2023
<u>AB</u>		<u>100MG</u>	<u>A204994 002</u>	Jan 05, 2023
<u>AB</u>		<u>150MG</u>	<u>A204994 003</u>	Jan 05, 2023
<u>AB</u>		<u>200MG</u>	<u>A204994 004</u>	Jan 05, 2023
<u>AB</u>	GLENMARK PHARMS LTD	<u>50MG</u>	<u>A205006 001</u>	Mar 17, 2022
<u>AB</u>		<u>100MG</u>	<u>A205006 002</u>	Mar 17, 2022
<u>AB</u>		<u>150MG</u>	<u>A205006 003</u>	Mar 17, 2022
<u>AB</u>		<u>200MG</u>	<u>A205006 004</u>	Mar 17, 2022
<u>AB</u>	HETERO LABS LTD V	<u>50MG</u>	<u>A204787 001</u>	Mar 17, 2022
<u>AB</u>		<u>100MG</u>	<u>A204787 002</u>	Mar 17, 2022
<u>AB</u>		<u>150MG</u>	<u>A204787 003</u>	Mar 17, 2022
<u>AB</u>		<u>200MG</u>	<u>A204787 004</u>	Mar 17, 2022
<u>AB</u>	INDOCO	<u>50MG</u>	<u>A208308 001</u>	Mar 17, 2022
<u>AB</u>		<u>100MG</u>	<u>A208308 002</u>	Mar 17, 2022
<u>AB</u>		<u>150MG</u>	<u>A208308 003</u>	Mar 17, 2022
<u>AB</u>		<u>200MG</u>	<u>A208308 004</u>	Mar 17, 2022
<u>AB</u>	KANCHAN HLTHCARE	<u>50MG</u>	<u>A218014 001</u>	Apr 02, 2024
<u>AB</u>		<u>100MG</u>	<u>A218014 002</u>	Apr 02, 2024
<u>AB</u>		<u>150MG</u>	<u>A218014 003</u>	Apr 02, 2024
<u>AB</u>		<u>200MG</u>	<u>A218014 004</u>	Apr 02, 2024
<u>AB</u>	MACLEODS PHARMS LTD	<u>50MG</u>	<u>A208466 001</u>	May 24, 2024
<u>AB</u>		<u>100MG</u>	<u>A208466 002</u>	May 24, 2024
<u>AB</u>		<u>150MG</u>	<u>A208466 003</u>	May 24, 2024
<u>AB</u>		<u>200MG</u>	<u>A208466 004</u>	May 24, 2024
<u>AB</u>	MSN LABS PVT LTD	<u>50MG</u>	<u>A204921 001</u>	Mar 17, 2022
<u>AB</u>		<u>100MG</u>	<u>A204921 002</u>	Mar 17, 2022
<u>AB</u>		<u>150MG</u>	<u>A204921 003</u>	Mar 17, 2022
<u>AB</u>		<u>200MG</u>	<u>A204921 004</u>	Mar 17, 2022
<u>AB</u>	SCIEGEN PHARMS INC	<u>50MG</u>	<u>A205237 001</u>	Mar 17, 2022
<u>AB</u>		<u>100MG</u>	<u>A205237 002</u>	Mar 17, 2022
<u>AB</u>		<u>150MG</u>	<u>A205237 003</u>	Mar 17, 2022
<u>AB</u>		<u>200MG</u>	<u>A205237 004</u>	Mar 17, 2022
<u>AB</u>	SUN PHARM	<u>50MG</u>	<u>A205031 001</u>	Mar 17, 2022
<u>AB</u>		<u>100MG</u>	<u>A205031 002</u>	Mar 17, 2022
<u>AB</u>		<u>150MG</u>	<u>A205031 003</u>	Mar 17, 2022
<u>AB</u>		<u>200MG</u>	<u>A205031 004</u>	Mar 17, 2022
<u>AB</u>	UNICHEM	<u>50MG</u>	<u>A213109 001</u>	Sep 04, 2024
<u>AB</u>		<u>100MG</u>	<u>A213109 002</u>	Sep 04, 2024
<u>AB</u>		<u>150MG</u>	<u>A213109 003</u>	Sep 04, 2024
<u>AB</u>		<u>200MG</u>	<u>A213109 004</u>	Sep 04, 2024
<u>VIMPAT</u>				
<u>AB</u>	+	<u>50MG</u>	<u>N022253 001</u>	Oct 28, 2008
<u>AB</u>	+	<u>100MG</u>	<u>N022253 002</u>	Oct 28, 2008
<u>AB</u>	+	<u>150MG</u>	<u>N022253 003</u>	Oct 28, 2008
<u>AB</u>	+	<u>200MG</u>	<u>N022253 004</u>	Oct 28, 2008

LACTULOSE

FOR SOLUTION; ORAL

LACTULOSE

!	CUMBERLAND PHARMS	10GM/PACKET	A074712 001	Dec 10, 1997
!		20GM/PACKET	A074712 002	Dec 10, 1997

SOLUTION; ORAL

LACTULOSE

<u>AA</u>	APOZEAL PHARMS	<u>10GM/15ML</u>	<u>A207786 001</u>	Jun 11, 2018
<u>AA</u>	AUROBINDO PHARMA LTD	<u>10GM/15ML</u>	<u>A074602 001</u>	Nov 14, 1996
<u>AA</u>	CHARTWELL RX	<u>10GM/15ML</u>	<u>A209517 001</u>	Nov 23, 2018
<u>AA</u>	FRESENIUS KABI	<u>10GM/15ML</u>	<u>A090503 001</u>	Jan 25, 2012
<u>AA</u>	LANNETT CO INC	<u>10GM/15ML</u>	<u>A075993 001</u>	Jul 26, 2001
<u>AA</u>	! PHARM ASSOC	<u>10GM/15ML</u>	<u>A074623 001</u>	Jul 30, 1996
<u>AA</u>	TARO	<u>10GM/15ML</u>	<u>A218858 001</u>	Dec 13, 2024
<u>AA</u>	XTTRIUM LABS INC	<u>10GM/15ML</u>	<u>A075911 001</u>	Feb 21, 2002

PRESCRIPTION DRUG PRODUCT LIST

LACTULOSE

SOLUTION;ORAL, RECTAL

GENERLAC

AA	CHARTWELL RX	<u>10GM/15ML</u>	A074603 001	Oct 31, 1996
-----------	--------------	------------------	--------------------	--------------

LACTULOSE

AA	APOZEAL PHARMS	<u>10GM/15ML</u>	A203762 001	Mar 27, 2015
AA	BAJAJ	<u>10GM/15ML</u>	A076645 001	Jul 28, 2003
AA	! FRESENIUS KABI	<u>10GM/15ML</u>	A090502 001	Jan 25, 2012

LAMIVUDINE

SOLUTION;ORAL

EPIVIR

AA	+! VIIV HLTHCARE	<u>10MG/ML</u>	N020596 001	Nov 17, 1995
-----------	------------------	----------------	--------------------	--------------

LAMIVUDINE

AA	AUROBINDO PHARMA	<u>10MG/ML</u>	A077695 001	Nov 21, 2016
AA	CHARTWELL MOLECULAR	<u>10MG/ML</u>	A203564 001	Oct 31, 2014
AA	HETERO LABS LTD III	<u>10MG/ML</u>	A091475 001	Oct 06, 2023

TABLET;ORAL

EPIVIR

AB	+ VIIV HLTHCARE	<u>150MG</u>	N020564 001	Nov 17, 1995
AB	+!	<u>300MG</u>	N020564 003	Jun 24, 2002

LAMIVUDINE

AB	ANNORA	<u>100MG</u>	A211306 001	Mar 21, 2019
AB	! APOTEX	<u>100MG</u>	A202941 001	Jan 02, 2014
AB		<u>150MG</u>	A091606 001	Dec 02, 2011
AB		<u>300MG</u>	A091606 002	Dec 02, 2011
AB	AUROBINDO PHARMA	<u>150MG</u>	A077464 001	Nov 21, 2016
AB		<u>300MG</u>	A077464 002	Nov 21, 2016
AB	BRECKENRIDGE	<u>150MG</u>	A203586 001	Nov 21, 2016
AB	CIPLA	<u>150MG</u>	A077221 001	Mar 03, 2017
AB		<u>300MG</u>	A077221 002	Mar 03, 2017
AB	HETERO LABS LTD V	<u>100MG</u>	A203260 001	Jan 02, 2014
AB		<u>150MG</u>	A203277 001	Jan 06, 2014
AB		<u>300MG</u>	A203277 002	Jan 06, 2014
AB	LUPIN LTD	<u>150MG</u>	A205217 001	Dec 18, 2014
AB		<u>300MG</u>	A205217 002	Dec 18, 2014
AB	MACLEODS PHARMS LTD	<u>150MG</u>	A090198 001	May 01, 2019
AB		<u>300MG</u>	A090198 002	May 01, 2019
AB	STRIDES PHARMA	<u>150MG</u>	A090457 001	Apr 19, 2018
AB		<u>300MG</u>	A090457 002	Apr 19, 2018
AB	UPSHER SMITH LABS	<u>150MG</u>	A206974 001	Nov 21, 2016
AB		<u>300MG</u>	A206974 002	Nov 21, 2016

LAMIVUDINE; TENOFOVIR DISOPROXIL FUMARATE

TABLET;ORAL

CIMDUO

+!	MYLAN LABS LTD	<u>300MG;300MG</u>	N022141 001	Feb 28, 2018
----	----------------	--------------------	--------------------	--------------

LAMIVUDINE; ZIDOVUDINE

TABLET;ORAL

LAMIVUDINE AND ZIDOVUDINE

AB	AUROBINDO PHARMA	<u>150MG;300MG</u>	A077558 001	May 05, 2017
AB	CIPLA	<u>150MG;300MG</u>	A077411 001	Sep 07, 2018
AB	EPIC PHARMA LLC	<u>150MG;300MG</u>	A206375 001	Apr 10, 2018
AB	! HETERO LABS LTD III	<u>150MG;300MG</u>	A079124 001	Sep 17, 2015
AB	HETERO LABS LTD V	<u>150MG;300MG</u>	A203259 001	Feb 03, 2014
AB	LUPIN LTD	<u>150MG;300MG</u>	A090246 001	May 15, 2012
AB	MACLEODS PHARMS LTD	<u>150MG;300MG</u>	A090679 001	Aug 29, 2018
AB	STRIDES PHARMA	<u>150MG;300MG</u>	A079128 001	May 13, 2015

LAMOTRIGINE

TABLET;ORAL

LAMICTAL

AB	+! GLAXOSMITHKLINE LLC	<u>25MG</u>	N020241 005	Dec 27, 1994
AB	+	<u>100MG</u>	N020241 001	Dec 27, 1994
AB	+	<u>150MG</u>	N020241 002	Dec 27, 1994
AB	+	<u>200MG</u>	N020241 003	Dec 27, 1994

LAMOTRIGINE

AB	ALEMBIC PHARMS LTD	<u>25MG</u>	A090607 001	Jan 13, 2011
AB		<u>100MG</u>	A090607 002	Jan 13, 2011
AB		<u>150MG</u>	A090607 003	Jan 13, 2011
AB		<u>200MG</u>	A090607 004	Jan 13, 2011
AB	ALKEM LABS LTD	<u>25MG</u>	A200694 001	Jun 14, 2013
AB		<u>100MG</u>	A200694 002	Jun 14, 2013

PRESCRIPTION DRUG PRODUCT LIST

LAMOTRIGINE

TABLET; ORAL

LAMOTRIGINE

<u>AB</u>		<u>150MG</u>	<u>A200694 003</u>	Jun 14, 2013
<u>AB</u>		<u>200MG</u>	<u>A200694 004</u>	Jun 14, 2013
<u>AB</u>	AUROBINDO PHARMA	<u>25MG</u>	<u>A078956 001</u>	Jan 27, 2009
<u>AB</u>		<u>100MG</u>	<u>A078956 002</u>	Jan 27, 2009
<u>AB</u>		<u>150MG</u>	<u>A078956 003</u>	Jan 27, 2009
<u>AB</u>		<u>200MG</u>	<u>A078956 004</u>	Jan 27, 2009
<u>AB</u>	DR REDDYS LABS LTD	<u>25MG</u>	<u>A076708 001</u>	Jan 27, 2009
<u>AB</u>		<u>100MG</u>	<u>A076708 002</u>	Jan 27, 2009
<u>AB</u>		<u>150MG</u>	<u>A076708 003</u>	Jan 27, 2009
<u>AB</u>		<u>200MG</u>	<u>A076708 004</u>	Jan 27, 2009
<u>AB</u>	GLENMARK PHARMS LTD	<u>25MG</u>	<u>A090169 001</u>	May 04, 2012
<u>AB</u>		<u>100MG</u>	<u>A090169 002</u>	May 04, 2012
<u>AB</u>		<u>150MG</u>	<u>A090169 003</u>	May 04, 2012
<u>AB</u>		<u>200MG</u>	<u>A090169 004</u>	May 04, 2012
<u>AB</u>	IPCA LABS	<u>25MG</u>	<u>A204499 001</u>	Sep 26, 2024
<u>AB</u>		<u>100MG</u>	<u>A204499 002</u>	Sep 26, 2024
<u>AB</u>		<u>150MG</u>	<u>A204499 003</u>	Sep 26, 2024
<u>AB</u>		<u>200MG</u>	<u>A204499 004</u>	Sep 26, 2024
<u>AB</u>	JUBILANT CADISTA	<u>25MG</u>	<u>A079132 001</u>	Jan 27, 2009
<u>AB</u>		<u>100MG</u>	<u>A079132 002</u>	Jan 27, 2009
<u>AB</u>		<u>150MG</u>	<u>A079132 003</u>	Jan 27, 2009
<u>AB</u>		<u>200MG</u>	<u>A079132 004</u>	Jan 27, 2009
<u>AB</u>	LUPIN LTD	<u>25MG</u>	<u>A078691 001</u>	Jun 01, 2010
<u>AB</u>		<u>100MG</u>	<u>A078691 002</u>	Jun 01, 2010
<u>AB</u>		<u>150MG</u>	<u>A078691 003</u>	Jun 01, 2010
<u>AB</u>		<u>200MG</u>	<u>A078691 004</u>	Jun 01, 2010
<u>AB</u>	RUBICON	<u>25MG</u>	<u>A078625 001</u>	Jan 27, 2009
<u>AB</u>		<u>100MG</u>	<u>A078625 002</u>	Jan 27, 2009
<u>AB</u>		<u>150MG</u>	<u>A078625 003</u>	Jan 27, 2009
<u>AB</u>		<u>200MG</u>	<u>A078625 004</u>	Jan 27, 2009
<u>AB</u>	TARO PHARM INDS	<u>25MG</u>	<u>A078525 001</u>	Jan 27, 2009
<u>AB</u>		<u>100MG</u>	<u>A078525 002</u>	Jan 27, 2009
<u>AB</u>		<u>150MG</u>	<u>A078525 003</u>	Jan 27, 2009
<u>AB</u>		<u>200MG</u>	<u>A078525 004</u>	Jan 27, 2009
<u>AB</u>	TORRENT PHARMS	<u>25MG</u>	<u>A078947 001</u>	Jan 27, 2009
<u>AB</u>		<u>100MG</u>	<u>A078947 002</u>	Jan 27, 2009
<u>AB</u>		<u>150MG</u>	<u>A078947 003</u>	Jan 27, 2009
<u>AB</u>		<u>200MG</u>	<u>A078947 004</u>	Jan 27, 2009
<u>AB</u>	UNICHEM LABS LTD	<u>25MG</u>	<u>A090170 001</u>	Oct 06, 2011
<u>AB</u>		<u>100MG</u>	<u>A090170 002</u>	Oct 06, 2011
<u>AB</u>		<u>150MG</u>	<u>A090170 003</u>	Oct 06, 2011
<u>AB</u>		<u>200MG</u>	<u>A090170 004</u>	Oct 06, 2011
<u>AB</u>	ZYDUS PHARMS USA	<u>25MG</u>	<u>A077633 001</u>	Jan 27, 2009
<u>AB</u>		<u>100MG</u>	<u>A077633 003</u>	Jan 27, 2009
<u>AB</u>		<u>150MG</u>	<u>A077633 004</u>	Jan 27, 2009
<u>AB</u>		<u>200MG</u>	<u>A077633 005</u>	Jan 27, 2009
		50MG	A077633 002	Jan 27, 2009
		250MG	A077633 006	Jan 27, 2009

TABLET, EXTENDED RELEASE; ORAL

LAMICTAL XR

<u>AB</u>	+	GLAXOSMITHKLINE LLC	<u>25MG</u>	<u>N022115 001</u>	May 29, 2009
<u>AB</u>	+	!	<u>50MG</u>	<u>N022115 002</u>	May 29, 2009
<u>AB</u>	+		<u>100MG</u>	<u>N022115 003</u>	May 29, 2009
<u>AB</u>	+	!	<u>200MG</u>	<u>N022115 004</u>	May 29, 2009
<u>AB</u>	+		<u>250MG</u>	<u>N022115 006</u>	Jun 21, 2011
<u>AB</u>	+		<u>300MG</u>	<u>N022115 005</u>	Apr 14, 2010

LAMOTRIGINE

<u>AB</u>	ACTAVIS ELIZABETH	<u>100MG</u>	<u>A200672 003</u>	Oct 17, 2013
<u>AB</u>		<u>200MG</u>	<u>A200672 004</u>	Oct 17, 2013
<u>AB</u>		<u>25MG</u>	<u>A200672 001</u>	Oct 17, 2013
<u>AB</u>		<u>50MG</u>	<u>A200672 002</u>	Oct 17, 2013
<u>AB</u>		<u>250MG</u>	<u>A200672 006</u>	Nov 13, 2013
<u>AB</u>		<u>300MG</u>	<u>A200672 005</u>	Oct 17, 2013
<u>AB</u>	ALEMBIC	<u>200MG</u>	<u>A211821 001</u>	Oct 01, 2024
<u>AB</u>		<u>250MG</u>	<u>A211821 002</u>	Oct 01, 2024
<u>AB</u>		<u>300MG</u>	<u>A211821 003</u>	Oct 01, 2024
<u>AB</u>	AMNEAL PHARMS	<u>25MG</u>	<u>A207497 001</u>	Nov 30, 2018
<u>AB</u>		<u>50MG</u>	<u>A207497 002</u>	Nov 30, 2018
<u>AB</u>		<u>100MG</u>	<u>A207497 003</u>	Nov 30, 2018

PRESCRIPTION DRUG PRODUCT LIST

LAMOTRIGINE

TABLET, EXTENDED RELEASE;ORAL

LAMOTRIGINE

<u>AB</u>		<u>200MG</u>	<u>A207497 004</u>	Nov 30, 2018
<u>AB</u>		<u>250MG</u>	<u>A207497 005</u>	Nov 30, 2018
<u>AB</u>		<u>300MG</u>	<u>A207497 006</u>	Nov 30, 2018
<u>AB</u>	DR REDDYS LABS LTD	<u>25MG</u>	<u>A202383 001</u>	Jun 19, 2013
<u>AB</u>		<u>50MG</u>	<u>A202383 002</u>	Jun 19, 2013
<u>AB</u>		<u>100MG</u>	<u>A202383 003</u>	Jun 19, 2013
<u>AB</u>		<u>200MG</u>	<u>A202383 004</u>	Jun 19, 2013
<u>AB</u>		<u>250MG</u>	<u>A202383 006</u>	Sep 06, 2018
<u>AB</u>		<u>300MG</u>	<u>A202383 005</u>	Jun 19, 2013
<u>AB</u>	ENDO OPERATIONS	<u>25MG</u>	<u>A201791 001</u>	Jan 18, 2013
<u>AB</u>		<u>50MG</u>	<u>A201791 002</u>	Jan 18, 2013
<u>AB</u>		<u>100MG</u>	<u>A201791 003</u>	Jan 18, 2013
<u>AB</u>		<u>200MG</u>	<u>A201791 004</u>	Jan 18, 2013
<u>AB</u>		<u>250MG</u>	<u>A201791 005</u>	Jan 18, 2013
<u>AB</u>		<u>300MG</u>	<u>A201791 006</u>	Jan 18, 2013
<u>AB</u>	RUBICON	<u>100MG</u>	<u>A202887 003</u>	Jun 17, 2013
<u>AB</u>		<u>200MG</u>	<u>A202887 004</u>	Jun 17, 2013
<u>AB</u>	SCIEGEN PHARMS INC	<u>25MG</u>	<u>A202498 001</u>	Jan 04, 2013
<u>AB</u>		<u>50MG</u>	<u>A202498 002</u>	Jan 04, 2013
<u>AB</u>		<u>100MG</u>	<u>A202498 003</u>	Jan 04, 2013
<u>AB</u>		<u>200MG</u>	<u>A202498 004</u>	Jan 04, 2013
<u>AB</u>		<u>300MG</u>	<u>A202498 005</u>	Jan 04, 2013
<u>AB</u>	TORRENT	<u>25MG</u>	<u>A203370 001</u>	Dec 23, 2013
<u>AB</u>		<u>50MG</u>	<u>A203370 002</u>	Dec 23, 2013
<u>AB</u>		<u>100MG</u>	<u>A203370 003</u>	Dec 23, 2013
<u>AB</u>	YILING	<u>25MG</u>	<u>A213949 001</u>	Dec 08, 2021
<u>AB</u>		<u>50MG</u>	<u>A213949 002</u>	Dec 08, 2021
<u>AB</u>		<u>100MG</u>	<u>A213949 003</u>	Dec 08, 2021
<u>AB</u>		<u>200MG</u>	<u>A213949 004</u>	Dec 08, 2021
<u>AB</u>		<u>250MG</u>	<u>A213949 005</u>	Dec 08, 2021
<u>AB</u>		<u>300MG</u>	<u>A213949 006</u>	Dec 08, 2021
<u>AB</u>	ZYDUS PHARMS	<u>25MG</u>	<u>A207763 001</u>	Apr 01, 2020
<u>AB</u>		<u>50MG</u>	<u>A207763 002</u>	Apr 01, 2020
<u>AB</u>		<u>100MG</u>	<u>A207763 003</u>	Apr 01, 2020
<u>AB</u>		<u>200MG</u>	<u>A207763 004</u>	Apr 01, 2020
<u>AB</u>		<u>250MG</u>	<u>A207763 005</u>	Apr 01, 2020
<u>AB</u>		<u>300MG</u>	<u>A207763 006</u>	Apr 01, 2020

TABLET, FOR SUSPENSION;ORAL

LAMICTAL CD

<u>AB</u>	+	GLAXOSMITHKLINE LLC	<u>2MG</u>	<u>N020764 004</u>	Sep 08, 2000
<u>AB</u>	+		<u>5MG</u>	<u>N020764 001</u>	Aug 24, 1998
<u>AB</u>	+	!	<u>25MG</u>	<u>N020764 002</u>	Aug 24, 1998

LAMOTRIGINE

<u>AB</u>	ALEMBIC	<u>5MG</u>	<u>A201168 001</u>	Jun 12, 2014
<u>AB</u>		<u>25MG</u>	<u>A201168 002</u>	Jun 12, 2014
<u>AB</u>	AUROBINDO PHARMA	<u>5MG</u>	<u>A090401 002</u>	Nov 04, 2009
<u>AB</u>		<u>25MG</u>	<u>A090401 003</u>	Nov 04, 2009
<u>AB</u>	DR REDDYS LABS LTD	<u>5MG</u>	<u>A076701 001</u>	Jan 22, 2009
<u>AB</u>		<u>25MG</u>	<u>A076701 002</u>	Jan 22, 2009
<u>AB</u>	GLENMARK PHARMS LTD	<u>5MG</u>	<u>A079099 001</u>	Feb 19, 2009
<u>AB</u>		<u>25MG</u>	<u>A079099 002</u>	Feb 19, 2009
<u>AB</u>	TARO	<u>5MG</u>	<u>A079204 001</u>	Feb 04, 2009
<u>AB</u>		<u>25MG</u>	<u>A079204 002</u>	Feb 04, 2009
<u>AB</u>	WATSON LABS	<u>2MG</u>	<u>A076928 001</u>	Jan 22, 2009
<u>AB</u>		<u>5MG</u>	<u>A076928 002</u>	Jan 22, 2009
<u>AB</u>		<u>25MG</u>	<u>A076928 003</u>	Jan 22, 2009
<u>AB</u>	ZYDUS PHARMS USA INC	<u>5MG</u>	<u>A078009 002</u>	Jan 22, 2009
<u>AB</u>		<u>25MG</u>	<u>A078009 003</u>	Jan 22, 2009

TABLET, ORALLY DISINTEGRATING;ORAL

LAMICTAL ODT

<u>AB</u>	+	GLAXOSMITHKLINE LLC	<u>25MG</u>	<u>N022251 001</u>	May 08, 2009
<u>AB</u>	+	!	<u>50MG</u>	<u>N022251 002</u>	May 08, 2009
<u>AB</u>	+		<u>100MG</u>	<u>N022251 003</u>	May 08, 2009
<u>AB</u>	+		<u>200MG</u>	<u>N022251 004</u>	May 08, 2009

LAMOTRIGINE

<u>AB</u>	AJANTA PHARMA LTD	<u>25MG</u>	<u>A213271 001</u>	Jan 19, 2021
<u>AB</u>		<u>50MG</u>	<u>A213271 002</u>	Jan 19, 2021
<u>AB</u>		<u>100MG</u>	<u>A213271 003</u>	Jan 19, 2021
<u>AB</u>		<u>200MG</u>	<u>A213271 004</u>	Jan 19, 2021

PRESCRIPTION DRUG PRODUCT LIST

LAMOTRIGINE

TABLET, ORALLY DISINTEGRATING;ORAL

LAMOTRIGINE

<u>AB</u>	AMRING PHARMS	<u>25MG</u>	<u>A214124 001</u>	Feb 03, 2022
<u>AB</u>		<u>50MG</u>	<u>A214124 002</u>	Feb 03, 2022
<u>AB</u>		<u>100MG</u>	<u>A214124 003</u>	Feb 03, 2022
<u>AB</u>		<u>200MG</u>	<u>A214124 004</u>	Feb 03, 2022
<u>AB</u>	ENDO OPERATIONS	<u>25MG</u>	<u>A204158 001</u>	Oct 27, 2015
<u>AB</u>		<u>50MG</u>	<u>A204158 002</u>	Oct 27, 2015
<u>AB</u>		<u>100MG</u>	<u>A204158 003</u>	Oct 27, 2015
<u>AB</u>		<u>200MG</u>	<u>A204158 004</u>	Oct 27, 2015
<u>AB</u>	IMPAX LABS INC	<u>25MG</u>	<u>A200828 001</u>	Jul 15, 2013
<u>AB</u>		<u>50MG</u>	<u>A200828 002</u>	Jul 15, 2013
<u>AB</u>		<u>100MG</u>	<u>A200828 003</u>	Jul 15, 2013
<u>AB</u>		<u>200MG</u>	<u>A200828 004</u>	Jul 15, 2013
<u>AB</u>	SCIEGEN PHARMS INC	<u>25MG</u>	<u>A206382 001</u>	Jun 17, 2016
<u>AB</u>		<u>50MG</u>	<u>A206382 002</u>	Jun 17, 2016
<u>AB</u>		<u>100MG</u>	<u>A206382 003</u>	Jun 17, 2016
<u>AB</u>		<u>200MG</u>	<u>A206382 004</u>	Jun 17, 2016

LANDIOLOL HYDROCHLORIDE

POWDER; INTRAVENOUS

RAPIBLYK

+! AOP ORPHAN

EQ 280MG BASE/VIAL

N217202 001 Nov 22, 2024

LANREOTIDE ACETATE

SOLUTION; SUBCUTANEOUS

LANREOTIDE ACETATE

<u>AB</u>	INVAGEN PHARMS	<u>EQ 60MG BASE/0.2ML (EQ 60MG BASE/0.2ML)</u>	<u>A217193 001</u>	May 21, 2024
<u>AB</u>		<u>EQ 90MG BASE/0.3ML (EQ 90MG BASE/0.3ML)</u>	<u>A217193 002</u>	May 21, 2024
<u>AB</u>		<u>EQ 120MG BASE/0.5ML (EQ 120MG BASE/0.5ML)</u>	<u>A217193 003</u>	May 21, 2024

SOMATULINE DEPOT

<u>AB</u>	+! IPSEN PHARMA	<u>EQ 60MG BASE/0.2ML (EQ 60MG BASE/0.2ML)</u>	<u>N022074 001</u>	Aug 30, 2007
<u>AB</u>	+!	<u>EQ 90MG BASE/0.3ML (EQ 90MG BASE/0.3ML)</u>	<u>N022074 002</u>	Aug 30, 2007
<u>AB</u>	+!	<u>EQ 120MG BASE/0.5ML (EQ 120MG BASE/0.5ML)</u>	<u>N022074 003</u>	Aug 30, 2007

LANREOTIDE ACETATE

+! INVAGEN PHARMS

EQ 60MG BASE/0.2ML (EQ 60MG BASE/0.2ML)

N215395 001 Dec 17, 2021

+!

EQ 90MG BASE/0.3ML (EQ 90MG BASE/0.3ML)

N215395 002 Dec 17, 2021

+!

EQ 120MG BASE/0.5ML (EQ 120MG

N215395 003 Dec 17, 2021

BASE/0.5ML)

LANSOPRAZOLE

CAPSULE, DELAYED REL PELLETS;ORAL

LANSOPRAZOLE

<u>AB</u>	ALKEM LABS LTD	<u>15MG</u>	<u>A207394 001</u>	Jan 18, 2019
<u>AB</u>		<u>30MG</u>	<u>A207394 002</u>	Jan 18, 2019
<u>AB</u>	CHARTWELL MOLECULAR	<u>15MG</u>	<u>A207156 001</u>	Sep 28, 2017
<u>AB</u>		<u>30MG</u>	<u>A207156 002</u>	Sep 28, 2017
<u>AB</u>	DR REDDYS LABS LTD	<u>15MG</u>	<u>A091269 001</u>	Oct 15, 2010
<u>AB</u>		<u>30MG</u>	<u>A091269 002</u>	Oct 15, 2010
<u>AB</u>	HETERO LABS LTD III	<u>15MG</u>	<u>A203083 001</u>	May 18, 2020
<u>AB</u>		<u>30MG</u>	<u>A203083 002</u>	May 18, 2020
<u>AB</u>	INVENTIA	<u>15MG</u>	<u>A205868 001</u>	Aug 30, 2017
<u>AB</u>		<u>30MG</u>	<u>A205868 002</u>	Aug 30, 2017
<u>AB</u>	MACLEODS PHARMS LTD	<u>15MG</u>	<u>A208671 001</u>	Nov 04, 2024
<u>AB</u>		<u>30MG</u>	<u>A208671 002</u>	Nov 04, 2024
<u>AB</u>	MYLAN PHARMS INC	<u>15MG</u>	<u>A090763 001</u>	Nov 10, 2009
<u>AB</u>		<u>30MG</u>	<u>A090763 002</u>	Nov 10, 2009
<u>AB</u>	NATCO PHARMA LTD	<u>15MG</u>	<u>A201921 001</u>	Dec 18, 2012
<u>AB</u>		<u>30MG</u>	<u>A201921 002</u>	Dec 18, 2012
<u>AB</u>	TEVA PHARMS	<u>15MG</u>	<u>A077255 001</u>	Nov 10, 2009
<u>AB</u>		<u>30MG</u>	<u>A077255 002</u>	Nov 10, 2009
<u>AB</u>	WOCKHARDT	<u>15MG</u>	<u>A202176 001</u>	Sep 14, 2012
<u>AB</u>		<u>30MG</u>	<u>A202176 002</u>	Sep 14, 2012
<u>AB</u>	XIROMED	<u>15MG</u>	<u>A203203 001</u>	Jul 25, 2016
<u>AB</u>		<u>30MG</u>	<u>A203203 002</u>	Jul 25, 2016
<u>AB</u>	ZYDUS HLTHCARE	<u>15MG</u>	<u>A202366 001</u>	Aug 19, 2013
<u>AB</u>		<u>30MG</u>	<u>A202366 002</u>	Aug 19, 2013

PREVACID

<u>AB</u>	+! TAKEDA PHARMS USA	<u>30MG</u>	<u>N020406 002</u>	May 10, 1995
-----------	----------------------	-------------	--------------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

LANSOPRAZOLE

TABLET, ORALLY DISINTEGRATING, DELAYED RELEASE;ORAL

LANSOPRAZOLE

AB	AUROBINDO PHARMA LTD	15MG	A207167 001	Mar 28, 2023
AB		30MG	A207167 002	Mar 28, 2023
AB	DR REDDYS	15MG	A210465 001	Feb 01, 2021
AB		30MG	A210465 002	Feb 01, 2021
AB	MYLAN	15MG	A202396 001	Nov 28, 2018
AB		30MG	A202396 002	Nov 28, 2018
AB	TEVA PHARMS USA	15MG	A208784 001	Sep 21, 2017
AB		30MG	A208784 002	Sep 21, 2017
AB	ZYDUS PHARMS	15MG	A200816 001	Nov 27, 2018
AB		30MG	A200816 002	Nov 27, 2018
<u>PREVACID</u>				
AB	+ TAKEDA PHARMS USA	15MG	N021428 001	Aug 30, 2002
AB	+!	30MG	N021428 002	Aug 30, 2002

LANTHANUM CARBONATE

POWDER;ORAL

FOSRENOL

	+ TAKEDA PHARMS USA	EQ 750MG BASE	N204734 001	Sep 24, 2014
	+!	EQ 1GM BASE	N204734 002	Sep 24, 2014

TABLET, CHEWABLE;ORAL

FOSRENOL

AB	+ TAKEDA PHARMS USA	EQ 500MG BASE	N021468 002	Oct 26, 2004
AB	+	EQ 750MG BASE	N021468 003	Nov 23, 2005
AB	+!	EQ 1GM BASE	N021468 004	Nov 23, 2005

LANTHANUM CARBONATE

AB	BARR	EQ 500MG BASE	A090977 001	Jan 27, 2022
AB		EQ 750MG BASE	A090977 002	Jan 27, 2022
AB		EQ 1GM BASE	A090977 003	Jan 27, 2022
AB	INVAGEN PHARMS	EQ 500MG BASE	A206868 001	Jan 24, 2022
AB		EQ 750MG BASE	A206868 002	Jan 24, 2022
AB		EQ 1GM BASE	A206868 003	Jan 24, 2022
AB	NATCO PHARMA LTD	EQ 500MG BASE	A090978 001	Aug 11, 2017
AB		EQ 750MG BASE	A090978 002	Aug 11, 2017
AB		EQ 1GM BASE	A090978 003	Aug 11, 2017

LAPATINIB DITOSYLATE

TABLET;ORAL

LAPATINIB DITOSYLATE

AB	NATCO PHARMA LTD	EQ 250MG BASE	A203007 001	Sep 29, 2020
AB	TEVA PHARMS USA INC	EQ 250MG BASE	A217968 001	Aug 16, 2024
<u>TYKERB</u>				
AB	+! NOVARTIS	EQ 250MG BASE	N022059 001	Mar 13, 2007

LAROTRECTINIB SULFATE

CAPSULE;ORAL

VITRAKVI

	+ BAYER HLTHCARE	EQ 25MG BASE	N210861 001	Nov 26, 2018
	+!	EQ 100MG BASE	N210861 002	Nov 26, 2018

SOLUTION;ORAL

VITRAKVI

	+! BAYER HEALTHCARE	EQ 20MG BASE/ML	N211710 001	Nov 26, 2018
--	---------------------	-----------------	-------------	--------------

LASMIDITAN SUCCINATE

TABLET;ORAL

REYVOW

	+ ELI LILLY AND CO	EQ 50MG BASE	N211280 001	Jan 31, 2020
	+!	EQ 100MG BASE	N211280 002	Jan 31, 2020

LATANOPROST

SOLUTION/DROPS;OPHTHALMIC

LATANOPROST

AT	AMRING PHARMS	0.005%	A200925 001	Mar 22, 2011
AT	BAUSCH AND LOMB	0.005%	A201006 001	Mar 22, 2011
AT	FDC LTD	0.005%	A202442 001	Apr 22, 2016
AT	GLAND	0.005%	A218257 001	Nov 26, 2024
AT	SANDOZ	0.005%	A091449 001	Mar 22, 2011
AT	SOMERSET	0.005%	A201786 001	Mar 22, 2011
<u>XALATAN</u>				
AT	+! UPJOHN	0.005%	N020597 001	Jun 05, 1996
	IYUZEH			
	+! THEA PHARMA	0.005%	N216472 001	Dec 13, 2022

PRESCRIPTION DRUG PRODUCT LIST

LATANOPROST; NETARSUDIL DIMESYLATE

SOLUTION/DROPS;OPHTHALMIC

ROCKLATAN

+! ALCON LABS INC 0.005%;EQ 0.02% BASE N208259 001 Mar 12, 2019

LATANOPROSTENE BUNOD

SOLUTION/DROPS;OPHTHALMIC

VYZULTA

+! BAUSCH AND LOMB 0.024% N207795 001 Nov 02, 2017

LAZERTINIB MESYLATE

TABLET;ORAL

LAZCLUZE

+ JANSSEN BIOTECH EQ 80MG BASE N219008 001 Aug 19, 2024

+! EQ 240MG BASE N219008 002 Aug 19, 2024

LEDIPASVIR; SOFOSBUVIR

PELLETS;ORAL

HARVONI

+ GILEAD SCIENCES INC 33.75MG;150MG/PACKET N212477 001 Aug 28, 2019

+! 45MG;200MG/PACKET N212477 002 Aug 28, 2019

TABLET;ORAL

HARVONI

+ GILEAD SCIENCES INC 45MG;200MG N205834 002 Aug 28, 2019

+! 90MG;400MG N205834 001 Oct 10, 2014

LEFAMULIN ACETATE

SOLUTION;INTRAVENOUS

XENLETA

+! HONG KONG EQ 150MG BASE/15ML (EQ 10MG BASE/ML) N211673 001 Aug 19, 2019

TABLET;ORAL

XENLETA

+! HONG KONG EQ 600MG BASE N211672 001 Aug 19, 2019

LEFLUNOMIDE

TABLET;ORAL

ARAVA**AB** + SANOFI AVENTIS US **10MG** **N020905 001** Sep 10, 1998**AB** +! **20MG** **N020905 002** Sep 10, 1998**LEFLUNOMIDE****AB** ABHAI LLC **10MG** **A212453 001** Jun 03, 2019**AB** **20MG** **A212453 002** Jun 03, 2019**AB** AET PHARMA **10MG** **A213497 001** May 10, 2021**AB** **20MG** **A213497 002** May 10, 2021**AB** ALEMBIC PHARMS LTD **10MG** **A091369 001** Nov 21, 2011**AB** **20MG** **A091369 002** Nov 21, 2011**AB** APOTEX INC **10MG** **A077090 001** Sep 13, 2005**AB** **20MG** **A077090 002** Sep 13, 2005**AB** AUROBINDO PHARMA **10MG** **A213652 001** Mar 29, 2021**AB** **20MG** **A213652 002** Mar 29, 2021**AB** HERITAGE **10MG** **A077086 001** Sep 13, 2005**AB** **20MG** **A077086 002** Sep 13, 2005**AB** LUPIN LTD **10MG** **A211863 001** Feb 04, 2020**AB** **20MG** **A211863 002** Feb 04, 2020**AB** WANBANG BIOPHARMS **10MG** **A077087 001** Sep 13, 2005**AB** **20MG** **A077087 002** Sep 13, 2005**AB** ZYDUS LIFESCIENCES **10MG** **A212308 001** Apr 24, 2019**AB** **20MG** **A212308 002** Apr 24, 2019

ARAVA

+! SANOFI AVENTIS US 100MG N020905 003 Sep 10, 1998

LEMBOREXANT

TABLET;ORAL

DAYVIGO

+ EISAI INC 5MG N212028 001 Apr 07, 2020

+! 10MG N212028 002 Apr 07, 2020

LENACAPAVIR SODIUM

SOLUTION;SUBCUTANEOUS

SUNLENCA

+! GILEAD SCIENCES INC EQ 463.5MG BASE/1.5ML (EQ 309MG BASE/ML) N215973 001 Dec 22, 2022

TABLET;ORAL

SUNLENCA

+! GILEAD SCIENCES INC EQ 300MG BASE N215974 001 Dec 22, 2022

PRESCRIPTION DRUG PRODUCT LIST

LENALIDOMIDE

CAPSULE; ORAL

LENALIDOMIDE

<u>AB</u>	APOTEX	<u>2.5MG</u>	<u>A211022</u>	<u>005</u>	Mar 07, 2023
<u>AB</u>		<u>5MG</u>	<u>A211022</u>	<u>001</u>	Aug 30, 2022
<u>AB</u>		<u>10MG</u>	<u>A211022</u>	<u>002</u>	Aug 30, 2022
<u>AB</u>		<u>15MG</u>	<u>A211022</u>	<u>003</u>	Aug 30, 2022
<u>AB</u>		<u>20MG</u>	<u>A211022</u>	<u>006</u>	Mar 07, 2023
<u>AB</u>		<u>25MG</u>	<u>A211022</u>	<u>004</u>	Aug 30, 2022
<u>AB</u>	ARROW INTL	<u>2.5MG</u>	<u>A201452</u>	<u>005</u>	Mar 06, 2023
<u>AB</u>		<u>5MG</u>	<u>A201452</u>	<u>001</u>	May 21, 2021
<u>AB</u>		<u>10MG</u>	<u>A201452</u>	<u>002</u>	May 21, 2021
<u>AB</u>		<u>15MG</u>	<u>A201452</u>	<u>003</u>	May 21, 2021
<u>AB</u>		<u>20MG</u>	<u>A201452</u>	<u>006</u>	Mar 06, 2023
<u>AB</u>		<u>25MG</u>	<u>A201452</u>	<u>004</u>	May 21, 2021
<u>AB</u>	CIPLA	<u>2.5MG</u>	<u>A214618</u>	<u>001</u>	Mar 06, 2023
<u>AB</u>		<u>5MG</u>	<u>A210435</u>	<u>001</u>	Sep 06, 2022
<u>AB</u>		<u>10MG</u>	<u>A210435</u>	<u>002</u>	Sep 06, 2022
<u>AB</u>		<u>15MG</u>	<u>A210435</u>	<u>003</u>	Sep 06, 2022
<u>AB</u>		<u>20MG</u>	<u>A210435</u>	<u>005</u>	Mar 06, 2023
<u>AB</u>		<u>25MG</u>	<u>A210435</u>	<u>004</u>	Sep 06, 2022
<u>AB</u>	DR REDDYS	<u>2.5MG</u>	<u>A209348</u>	<u>001</u>	Oct 14, 2021
<u>AB</u>		<u>5MG</u>	<u>A209348</u>	<u>003</u>	Aug 30, 2022
<u>AB</u>		<u>10MG</u>	<u>A209348</u>	<u>004</u>	Aug 30, 2022
<u>AB</u>		<u>15MG</u>	<u>A209348</u>	<u>005</u>	Aug 30, 2022
<u>AB</u>		<u>20MG</u>	<u>A209348</u>	<u>002</u>	Oct 14, 2021
<u>AB</u>		<u>25MG</u>	<u>A209348</u>	<u>006</u>	Aug 30, 2022
<u>AB</u>	EUGIA PHARMA	<u>2.5MG</u>	<u>A213885</u>	<u>001</u>	Mar 06, 2023
<u>AB</u>		<u>5MG</u>	<u>A213885</u>	<u>002</u>	Mar 06, 2023
<u>AB</u>		<u>10MG</u>	<u>A213885</u>	<u>003</u>	Mar 06, 2023
<u>AB</u>		<u>15MG</u>	<u>A213885</u>	<u>004</u>	Mar 06, 2023
<u>AB</u>		<u>20MG</u>	<u>A213885</u>	<u>005</u>	Mar 06, 2023
<u>AB</u>		<u>25MG</u>	<u>A213885</u>	<u>006</u>	Mar 06, 2023
<u>AB</u>	HETERO LABS LTD V	<u>2.5MG</u>	<u>A212414</u>	<u>001</u>	May 11, 2023
<u>AB</u>		<u>5MG</u>	<u>A212414</u>	<u>002</u>	May 11, 2023
<u>AB</u>		<u>10MG</u>	<u>A212414</u>	<u>003</u>	May 11, 2023
<u>AB</u>		<u>15MG</u>	<u>A212414</u>	<u>004</u>	May 11, 2023
<u>AB</u>		<u>20MG</u>	<u>A212414</u>	<u>005</u>	May 11, 2023
<u>AB</u>		<u>25MG</u>	<u>A212414</u>	<u>006</u>	May 11, 2023
<u>AB</u>	LOTUS PHARM CO LTD	<u>2.5MG</u>	<u>A210480</u>	<u>005</u>	Mar 06, 2023
<u>AB</u>		<u>5MG</u>	<u>A210480</u>	<u>001</u>	Aug 31, 2022
<u>AB</u>		<u>10MG</u>	<u>A210480</u>	<u>002</u>	Aug 31, 2022
<u>AB</u>		<u>15MG</u>	<u>A210480</u>	<u>003</u>	Aug 31, 2022
<u>AB</u>		<u>20MG</u>	<u>A210480</u>	<u>006</u>	Mar 06, 2023
<u>AB</u>		<u>25MG</u>	<u>A210480</u>	<u>004</u>	Aug 31, 2022
<u>AB</u>	MYLAN	<u>2.5MG</u>	<u>A213912</u>	<u>005</u>	Mar 06, 2023
<u>AB</u>		<u>5MG</u>	<u>A213912</u>	<u>001</u>	Aug 30, 2022
<u>AB</u>		<u>10MG</u>	<u>A213912</u>	<u>002</u>	Aug 30, 2022
<u>AB</u>		<u>15MG</u>	<u>A213912</u>	<u>003</u>	Aug 30, 2022
<u>AB</u>		<u>20MG</u>	<u>A213912</u>	<u>006</u>	Mar 06, 2023
<u>AB</u>		<u>25MG</u>	<u>A213912</u>	<u>004</u>	Aug 30, 2022
<u>AB</u>	NOVUGEN	<u>2.5MG</u>	<u>A217281</u>	<u>001</u>	Sep 11, 2024
<u>AB</u>		<u>5MG</u>	<u>A217281</u>	<u>002</u>	Sep 11, 2024
<u>AB</u>		<u>10MG</u>	<u>A217281</u>	<u>003</u>	Sep 11, 2024
<u>AB</u>		<u>15MG</u>	<u>A217281</u>	<u>004</u>	Sep 11, 2024
<u>AB</u>		<u>20MG</u>	<u>A217281</u>	<u>005</u>	Sep 11, 2024
<u>AB</u>		<u>25MG</u>	<u>A217281</u>	<u>006</u>	Sep 11, 2024
<u>AB</u>	SUN PHARM	<u>2.5MG</u>	<u>A211846</u>	<u>005</u>	Mar 06, 2023
<u>AB</u>		<u>5MG</u>	<u>A211846</u>	<u>001</u>	Feb 08, 2023
<u>AB</u>		<u>10MG</u>	<u>A211846</u>	<u>002</u>	Feb 08, 2023
<u>AB</u>		<u>15MG</u>	<u>A211846</u>	<u>003</u>	Feb 08, 2023
<u>AB</u>		<u>20MG</u>	<u>A211846</u>	<u>006</u>	Mar 06, 2023
<u>AB</u>		<u>25MG</u>	<u>A211846</u>	<u>004</u>	Feb 08, 2023
<u>AB</u>	ZYDUS PHARMS	<u>2.5MG</u>	<u>A210154</u>	<u>005</u>	Mar 06, 2023
<u>AB</u>		<u>5MG</u>	<u>A210154</u>	<u>001</u>	Sep 12, 2022
<u>AB</u>		<u>10MG</u>	<u>A210154</u>	<u>002</u>	Sep 12, 2022
<u>AB</u>		<u>15MG</u>	<u>A210154</u>	<u>003</u>	Sep 12, 2022
<u>AB</u>		<u>20MG</u>	<u>A210154</u>	<u>006</u>	Mar 06, 2023
<u>AB</u>		<u>25MG</u>	<u>A210154</u>	<u>004</u>	Sep 12, 2022
<u>REVLIMID</u>					
<u>AB</u>	+ BRISTOL MYERS SQUIBB	<u>2.5MG</u>	<u>N021880</u>	<u>005</u>	Dec 21, 2011
<u>AB</u>	+	<u>5MG</u>	<u>N021880</u>	<u>001</u>	Dec 27, 2005

PRESCRIPTION DRUG PRODUCT LIST

LENALIDOMIDE

CAPSULE; ORAL

REVLIMID

<u>AB</u>	+		<u>10MG</u>	<u>N021880</u>	<u>002</u>	Dec 27, 2005
<u>AB</u>	+		<u>15MG</u>	<u>N021880</u>	<u>003</u>	Jun 29, 2006
<u>AB</u>	+		<u>20MG</u>	<u>N021880</u>	<u>006</u>	Jun 05, 2013
<u>AB</u>	+	!	<u>25MG</u>	<u>N021880</u>	<u>004</u>	Jun 29, 2006

LENIOLISIB PHOSPHATE

TABLET; ORAL

JOENJA

+	!	PHARMING	EQ 70MG BASE	N217759	001	Mar 24, 2023
---	---	----------	--------------	---------	-----	--------------

LENVATINIB MESYLATE

CAPSULE; ORAL

LENVIMA

+		EISAI INC	EQ 4MG BASE	N206947	001	Feb 13, 2015
---	--	-----------	-------------	---------	-----	--------------

+	!		EQ 10MG BASE	N206947	002	Feb 13, 2015
---	---	--	--------------	---------	-----	--------------

LETERMOVIR

PELLETS; ORAL

PREVMIS

+		MSD	20MG/PACKET	N219104	001	Aug 30, 2024
---	--	-----	-------------	---------	-----	--------------

+	!		120MG/PACKET	N219104	002	Aug 30, 2024
---	---	--	--------------	---------	-----	--------------

SOLUTION; INTRAVENOUS

PREVMIS

+	!	MERCK SHARP DOHME	240MG/12ML (20MG/ML)	N209940	001	Nov 08, 2017
---	---	-------------------	----------------------	---------	-----	--------------

+	!		480MG/24ML (20MG/ML)	N209940	002	Nov 08, 2017
---	---	--	----------------------	---------	-----	--------------

TABLET; ORAL

PREVMIS

+		MERCK SHARP DOHME	240MG	N209939	001	Nov 08, 2017
---	--	-------------------	-------	---------	-----	--------------

+	!		480MG	N209939	002	Nov 08, 2017
---	---	--	-------	---------	-----	--------------

LETROZOLE

TABLET; ORAL

FEMARA

<u>AB</u>	+	!	NOVARTIS PHARMS	<u>2.5MG</u>	<u>N020726</u>	<u>001</u>	Jul 25, 1997
-----------	---	---	-----------------	--------------	----------------	------------	--------------

LETROZOLE

<u>AB</u>			ACCORD HLTHCARE	<u>2.5MG</u>	<u>A090934</u>	<u>001</u>	Jun 03, 2011
<u>AB</u>			BEIJING YILING	<u>2.5MG</u>	<u>A205869</u>	<u>001</u>	Nov 14, 2018
<u>AB</u>			EUGIA PHARMA	<u>2.5MG</u>	<u>A211717</u>	<u>001</u>	Jan 11, 2019
<u>AB</u>			NATCO PHARMA LTD	<u>2.5MG</u>	<u>A200161</u>	<u>001</u>	Jun 03, 2011
<u>AB</u>			TEVA PHARMS	<u>2.5MG</u>	<u>A090289</u>	<u>001</u>	Jun 03, 2011

LETROZOLE; RIBOCICLIB SUCCINATE

TABLET; ORAL

KISQALI FEMARA CO-PACK (COPACKAGED)

+	!	NOVARTIS	2.5MG;EQ 200MG BASE	N209935	001	May 04, 2017
---	---	----------	---------------------	---------	-----	--------------

LEUCOVORIN CALCIUM

INJECTABLE; INJECTION

LEUCOVORIN CALCIUM

<u>AP</u>			HAINAN POLY	<u>EQ 50MG BASE/VIAL</u>	<u>A217021</u>	<u>001</u>	Jul 10, 2023
<u>AP</u>				<u>EQ 100MG BASE/VIAL</u>	<u>A217021</u>	<u>002</u>	Jul 10, 2023
<u>AP</u>				<u>EQ 200MG BASE/VIAL</u>	<u>A217021</u>	<u>003</u>	Jul 10, 2023
<u>AP</u>				<u>EQ 350MG BASE/VIAL</u>	<u>A217021</u>	<u>005</u>	Dec 19, 2024
<u>AP</u>				<u>EQ 500MG BASE/VIAL</u>	<u>A217021</u>	<u>004</u>	Jul 10, 2023
<u>AP</u>	!		HIKMA	<u>EQ 50MG BASE/VIAL</u>	<u>A089384</u>	<u>001</u>	Sep 14, 1987
<u>AP</u>	!			<u>EQ 100MG BASE/VIAL</u>	<u>A089717</u>	<u>001</u>	Mar 28, 1988
<u>AP</u>			MEITHEAL	<u>EQ 50MG BASE/VIAL</u>	<u>A216590</u>	<u>001</u>	Jul 19, 2023
<u>AP</u>				<u>EQ 100MG BASE/VIAL</u>	<u>A216590</u>	<u>002</u>	Jul 19, 2023
<u>AP</u>				<u>EQ 200MG BASE/VIAL</u>	<u>A216590</u>	<u>003</u>	Jul 19, 2023
<u>AP</u>				<u>EQ 350MG BASE/VIAL</u>	<u>A216590</u>	<u>004</u>	Jul 19, 2023
<u>AP</u>				<u>EQ 500MG BASE/VIAL</u>	<u>A216590</u>	<u>005</u>	Jul 19, 2023
				<u>LEUCOVORIN CALCIUM PRESERVATIVE FREE</u>			
<u>AP</u>			FRESENIUS KABI USA	<u>EQ 200MG BASE/VIAL</u>	<u>A040258</u>	<u>001</u>	Feb 26, 1999
<u>AP</u>	!			<u>EQ 500MG BASE/VIAL</u>	<u>A040286</u>	<u>001</u>	Feb 26, 1999
<u>AP</u>	!		HIKMA	<u>EQ 200MG BASE/VIAL</u>	<u>A040056</u>	<u>001</u>	May 23, 1995
<u>AP</u>	!			<u>EQ 350MG BASE/VIAL</u>	<u>A040335</u>	<u>001</u>	Apr 20, 2000
<u>AP</u>			MYLAN LABS LTD	<u>EQ 100MG BASE/VIAL</u>	<u>A203800</u>	<u>001</u>	May 19, 2017
<u>AP</u>				<u>EQ 200MG BASE/VIAL</u>	<u>A203800</u>	<u>002</u>	May 19, 2017
<u>AP</u>				<u>EQ 350MG BASE/VIAL</u>	<u>A203800</u>	<u>003</u>	May 19, 2017
<u>AP</u>			SAGENT PHARMS	<u>EQ 50MG BASE/VIAL</u>	<u>A200753</u>	<u>001</u>	Sep 06, 2012
<u>AP</u>				<u>EQ 100MG BASE/VIAL</u>	<u>A200753</u>	<u>002</u>	Sep 06, 2012
<u>AP</u>				<u>EQ 200MG BASE/VIAL</u>	<u>A200753</u>	<u>003</u>	Sep 06, 2012

PRESCRIPTION DRUG PRODUCT LIST

LEUCOVORIN CALCIUM

INJECTABLE; INJECTION

LEUCOVORIN CALCIUM PRESERVATIVE FREE

<u>AP</u>		<u>EQ 350MG BASE/VIAL</u>	<u>A200855 001</u>	Sep 06, 2012
<u>AP</u>	SAGENT PHARMS INC	<u>EQ 500MG BASE/VIAL</u>	<u>A209110 001</u>	Oct 26, 2017
	SOLUTION; INTRAMUSCULAR, INTRAVENOUS			
	LEUCOVORIN CALCIUM			
	! FRESenius KABI USA	EQ 100MG BASE/10ML (EQ 10MG BASE/ML)	A207241 001	Mar 14, 2018
	!	EQ 500MG BASE/50ML (EQ 10MG BASE/ML)	A207226 001	Jul 27, 2018

TABLET; ORAL

LEUCOVORIN CALCIUM

<u>AB</u>	BARR	<u>EQ 5MG BASE</u>	<u>A071198 001</u>	Sep 24, 1987
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>A071199 001</u>	Sep 24, 1987
<u>AB</u>	EPIC PHARMA LLC	<u>EQ 5MG BASE</u>	<u>A074544 001</u>	Aug 28, 1997
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A074544 003</u>	May 19, 2021
<u>AB</u>		<u>EQ 15MG BASE</u>	<u>A074544 004</u>	May 19, 2021
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>A074544 002</u>	Aug 28, 1997
<u>AB</u>	HIKMA	<u>EQ 5MG BASE</u>	<u>A072733 001</u>	Feb 22, 1993
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A072734 001</u>	Feb 22, 1993
<u>AB</u>		<u>EQ 15MG BASE</u>	<u>A072735 001</u>	Feb 22, 1993
<u>AB</u>	!	<u>EQ 25MG BASE</u>	<u>A072736 001</u>	Feb 22, 1993
<u>AB</u>	LEADING	<u>EQ 5MG BASE</u>	<u>A213929 001</u>	Oct 22, 2020
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A213929 002</u>	Oct 22, 2020
<u>AB</u>		<u>EQ 15MG BASE</u>	<u>A213929 003</u>	Oct 22, 2020
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>A213929 004</u>	Oct 22, 2020
<u>AB</u>	NOVAST LABS	<u>EQ 5MG BASE</u>	<u>A211132 001</u>	Jul 30, 2020
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A211132 002</u>	Jul 30, 2020
<u>AB</u>		<u>EQ 15MG BASE</u>	<u>A211132 003</u>	Jul 30, 2020
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>A211132 004</u>	Jul 30, 2020

LEUPROLIDE ACETATE

FOR SUSPENSION; INTRAMUSCULAR

LEUPROLIDE ACETATE FOR DEPOT SUSPENSION

+!	INVAGEN PHARMS	22.5MG/VIAL	N205054 001	Aug 28, 2018
----	----------------	-------------	-------------	--------------

INJECTABLE; INJECTION

LUPRON DEPOT

+!	ABBVIE ENDOCRINE INC	3.75MG	N020011 002	Oct 26, 1995
+!		7.5MG	N019732 001	Jan 26, 1989
+!		11.25MG	N020708 001	Mar 07, 1997
+		22.5MG	N020517 001	Dec 22, 1995
+!		30MG	N020517 002	May 30, 1997
+!		45MG	N020517 003	Jun 17, 2011

POWDER; INTRAMUSCULAR

LUPRON DEPOT-PED KIT

+!	ABBVIE ENDOCRINE INC	7.5MG	N020263 002	Apr 16, 1993
+!		11.25MG	N020263 005	Jan 21, 1994
+!		11.25MG	N020263 007	Aug 15, 2011
+!		15MG	N020263 006	Jan 21, 1994
+!		30MG	N020263 008	Aug 15, 2011
+!		45MG	N020263 009	Apr 14, 2023

POWDER; SUBCUTANEOUS

ELIGARD KIT

+!	TOLMAR	7.5MG	N021343 001	Jan 23, 2002
+!		22.5MG	N021379 001	Jul 24, 2002
+!		30MG	N021488 001	Feb 13, 2003
+!		45MG	N021731 001	Dec 14, 2004

FENSOLVI KIT

+!	TOLMAR	45MG	N213150 001	May 01, 2020
----	--------	------	-------------	--------------

SOLUTION; SUBCUTANEOUS

LEUPROLIDE ACETATE

<u>AP</u>	AMNEAL	<u>14MG/2.8ML (1MG/0.2ML)</u>	<u>A215336 001</u>	Oct 28, 2022
<u>AP</u>	EUGIA PHARMA	<u>14MG/2.8ML (1MG/0.2ML)</u>	<u>A212963 001</u>	Jun 06, 2022
<u>AP</u>	MEITHEAL	<u>14MG/2.8ML (1MG/0.2ML)</u>	<u>A075471 001</u>	Oct 25, 2000
<u>AP</u>	RK PHARMA	<u>14MG/2.8ML (1MG/0.2ML)</u>	<u>A213829 001</u>	Aug 13, 2021
<u>AP</u>	! SANDOZ	<u>14MG/2.8ML (1MG/0.2ML)</u>	<u>A074728 001</u>	Aug 04, 1998
<u>AP</u>	SUN PHARM	<u>14MG/2.8ML (1MG/0.2ML)</u>	<u>A078885 001</u>	Mar 09, 2009
<u>AP</u>	UBI	<u>14MG/2.8ML (1MG/0.2ML)</u>	<u>A217957 001</u>	Oct 17, 2024

PRESCRIPTION DRUG PRODUCT LIST

LEUPROLIDE MESYLATE

EMULSION; SUBCUTANEOUS

CAMCEVI KIT

+! ACCORD

EQ 42MG BASE

N211488 001 May 25, 2021

LEVACETYLLAUCINE

FOR SUSPENSION; ORAL

AQNEURSA

+! INTRABIO

1GM/PACKET

N219132 001 Sep 24, 2024

LEVALBUTEROL HYDROCHLORIDE

SOLUTION; INHALATION

LEVALBUTEROL HYDROCHLORIDE

<u>AN</u>		CIPLA	<u>EQ 0.021% BASE</u>	<u>A078171 002</u>	Dec 13, 2013
<u>AN</u>			<u>EQ 0.042% BASE</u>	<u>A078171 003</u>	Dec 13, 2013
<u>AN</u>			<u>EQ 0.0103% BASE</u>	<u>A078171 001</u>	Dec 13, 2013
<u>AN</u>		IMPAX LABS INC	<u>EQ 0.0103% BASE</u>	<u>A077756 003</u>	Apr 09, 2008
<u>AN</u>			<u>EQ 0.021% BASE</u>	<u>A077756 001</u>	Apr 09, 2008
<u>AN</u>			<u>EQ 0.042% BASE</u>	<u>A077756 002</u>	Apr 09, 2008
<u>AN</u>		LUOXIN AUROVITAS	<u>EQ 0.25% BASE</u>	<u>A207628 001</u>	Jan 31, 2017
<u>AN</u>			<u>EQ 0.0103% BASE</u>	<u>A207625 001</u>	Dec 30, 2016
<u>AN</u>			<u>EQ 0.021% BASE</u>	<u>A207625 002</u>	Dec 30, 2016
<u>AN</u>			<u>EQ 0.042% BASE</u>	<u>A207625 003</u>	Dec 30, 2016
<u>AN</u>		MANKIND PHARMA	<u>EQ 0.0103% BASE</u>	<u>A218770 001</u>	Oct 21, 2024
<u>AN</u>			<u>EQ 0.021% BASE</u>	<u>A218770 002</u>	Oct 21, 2024
<u>AN</u>			<u>EQ 0.042% BASE</u>	<u>A218770 003</u>	Oct 21, 2024
<u>AN</u>	!	MYLAN SPECIALITY LP	<u>EQ 0.25% BASE</u>	<u>A078309 001</u>	Mar 20, 2009
<u>AN</u>		RITEDOSE CORP	<u>EQ 0.0103% BASE</u>	<u>A203653 001</u>	Mar 22, 2016
<u>AN</u>			<u>EQ 0.021% BASE</u>	<u>A203653 002</u>	Mar 22, 2016
<u>AN</u>			<u>EQ 0.042% BASE</u>	<u>A203653 003</u>	Mar 22, 2016
<u>AN</u>		TEVA PARENTERAL	<u>EQ 0.25% BASE</u>	<u>A200875 001</u>	Sep 11, 2014
<u>AN</u>	!	TEVA PHARMS USA	<u>EQ 0.0103% BASE</u>	<u>A090297 001</u>	Apr 26, 2013
<u>AN</u>	!		<u>EQ 0.021% BASE</u>	<u>A090297 002</u>	Apr 26, 2013
<u>AN</u>	!		<u>EQ 0.042% BASE</u>	<u>A090297 003</u>	Apr 26, 2013

LEVALBUTEROL TARTRATE

AEROSOL, METERED; INHALATION

XOPENEX HFA

+! LUPIN

EQ 0.045MG BASE/INH

N021730 001 Mar 11, 2005

LEVETIRACETAM

INJECTABLE; INTRAVENOUS

KEPPRA

<u>AP</u>	+	UCB INC	<u>500MG/5ML (100MG/ML)</u>	<u>N021872 001</u>	Jul 31, 2006
<u>AP</u>		EUGIA PHARMA	<u>500MG/5ML (100MG/ML)</u>	<u>A204312 001</u>	Feb 01, 2016
<u>AP</u>		FRESENIUS KABI USA	<u>500MG/5ML (100MG/ML)</u>	<u>A090876 001</u>	Aug 13, 2015
<u>AP</u>		HAINAN POLY PHARM	<u>500MG/5ML (100MG/ML)</u>	<u>A209781 001</u>	Mar 20, 2018
<u>AP</u>		HIKMA FARMACEUTICA	<u>500MG/5ML (100MG/ML)</u>	<u>A090981 001</u>	Oct 13, 2011
<u>AP</u>		HOSPIRA INC	<u>500MG/5ML (100MG/ML)</u>	<u>A202869 001</u>	Apr 06, 2012
<u>AP</u>		MICRO LABS	<u>500MG/5ML (100MG/ML)</u>	<u>A211954 001</u>	Aug 09, 2019
<u>AP</u>		MSN	<u>500MG/5ML (100MG/ML)</u>	<u>A215980 001</u>	Nov 15, 2022
<u>AP</u>		MYLAN LABS LTD	<u>500MG/5ML (100MG/ML)</u>	<u>A203308 001</u>	Sep 16, 2016
<u>AP</u>		PRINSON INC	<u>500MG/5ML (100MG/ML)</u>	<u>A209474 001</u>	Mar 28, 2022
<u>AP</u>		SAGENT PHARMS	<u>500MG/5ML (100MG/ML)</u>	<u>A091627 001</u>	Jun 26, 2013
<u>AP</u>		XGEN PHARMS	<u>500MG/5ML (100MG/ML)</u>	<u>A091485 001</u>	Aug 05, 2011

LEVETIRACETAM IN SODIUM CHLORIDE

<u>AP</u>		B BRAUN MEDICAL INC	<u>500MG/100ML (5MG/ML)</u>	<u>A209705 001</u>	Feb 27, 2024
<u>AP</u>			<u>1GM/100ML (10MG/ML)</u>	<u>A209705 002</u>	Feb 27, 2024
<u>AP</u>			<u>1.5GM/100ML (15MG/ML)</u>	<u>A209705 003</u>	Feb 27, 2024
<u>AP</u>		BAXTER HLTHCARE CORP	<u>500MG/100ML (5MG/ML)</u>	<u>A217059 001</u>	Oct 02, 2023
<u>AP</u>			<u>1GM/100ML (10MG/ML)</u>	<u>A217059 002</u>	Oct 02, 2023
<u>AP</u>			<u>1.5GM/100ML (15MG/ML)</u>	<u>A217059 003</u>	Oct 02, 2023
<u>AP</u>		EUGIA PHARMA	<u>500MG/100ML (5MG/ML)</u>	<u>A207160 001</u>	Jan 04, 2017
<u>AP</u>			<u>1GM/100ML (10MG/ML)</u>	<u>A207160 002</u>	Jan 04, 2017
<u>AP</u>			<u>1.5GM/100ML (15MG/ML)</u>	<u>A207160 003</u>	Jan 04, 2017
<u>AP</u>		GLAND PHARMA LTD	<u>500MG/100ML (5MG/ML)</u>	<u>A206880 001</u>	Oct 25, 2017
<u>AP</u>			<u>1GM/100ML (10MG/ML)</u>	<u>A206880 002</u>	Oct 25, 2017
<u>AP</u>			<u>1.5GM/100ML (15MG/ML)</u>	<u>A206880 003</u>	Oct 25, 2017
<u>AP</u>		HIKMA	<u>500MG/100ML (5MG/ML)</u>	<u>A211356 001</u>	Feb 12, 2024
<u>AP</u>			<u>1GM/100ML (10MG/ML)</u>	<u>A211356 002</u>	Feb 12, 2024
<u>AP</u>			<u>1.5GM/100ML (15MG/ML)</u>	<u>A211356 003</u>	Feb 12, 2024
<u>AP</u>	+	HQ SPCLT PHARMA	<u>500MG/100ML (5MG/ML)</u>	<u>N202543 001</u>	Nov 09, 2011
<u>AP</u>	+		<u>1GM/100ML (10MG/ML)</u>	<u>N202543 002</u>	Nov 09, 2011

PRESCRIPTION DRUG PRODUCT LIST

LEVETIRACETAM

INJECTABLE; INTRAVENOUS

LEVETIRACETAM IN SODIUM CHLORIDE

<u>AP</u>	<u>+!</u>	<u>1.5GM/100ML (15MG/ML)</u>	<u>N202543</u>	<u>003</u>	Nov 09, 2011
<u>AP</u>	NEXUS	<u>500MG/100ML (5MG/ML)</u>	<u>A213532</u>	<u>001</u>	Jul 06, 2020
<u>AP</u>		<u>1GM/100ML (10MG/ML)</u>	<u>A213532</u>	<u>002</u>	Jul 06, 2020
<u>AP</u>		<u>1.5GM/100ML (15MG/ML)</u>	<u>A213532</u>	<u>003</u>	Jul 06, 2020

SOLUTION; ORAL

KEPPRA

<u>AA</u>	<u>+!</u>	UCB INC	<u>100MG/ML</u>	<u>N021505</u>	<u>001</u>	Jul 15, 2003
-----------	-----------	---------	-----------------	----------------	------------	--------------

LEVETIRACETAM

<u>AA</u>		ACTAVIS MID ATLANTIC	<u>100MG/ML</u>	<u>A078976</u>	<u>001</u>	Jan 15, 2009
<u>AA</u>		ALEMBIC	<u>100MG/ML</u>	<u>A203067</u>	<u>001</u>	May 09, 2013
<u>AA</u>		AMNEAL PHARMS	<u>100MG/ML</u>	<u>A090992</u>	<u>001</u>	Oct 27, 2009
<u>AA</u>		AUROBINDO PHARMA	<u>100MG/ML</u>	<u>A079063</u>	<u>001</u>	Jan 15, 2009
<u>AA</u>		BELCHER	<u>100MG/ML</u>	<u>A090461</u>	<u>001</u>	Sep 30, 2010
<u>AA</u>		CHARTWELL MOLECULAR	<u>100MG/ML</u>	<u>A090263</u>	<u>001</u>	Apr 03, 2009
<u>AA</u>		HETERO LABS LTD III	<u>100MG/ML</u>	<u>A203052</u>	<u>001</u>	Feb 28, 2013
<u>AA</u>		LUPIN LTD	<u>100MG/ML</u>	<u>A090893</u>	<u>001</u>	Oct 17, 2011
<u>AA</u>		MSN	<u>100MG/ML</u>	<u>A214757</u>	<u>001</u>	Jul 15, 2022
<u>AA</u>		PHARM ASSOC	<u>100MG/ML</u>	<u>A201157</u>	<u>001</u>	Jun 04, 2015
<u>AA</u>		QUAGEN	<u>100MG/ML</u>	<u>A090079</u>	<u>001</u>	Apr 11, 2012
<u>AA</u>		STRIDES PHARMA	<u>100MG/ML</u>	<u>A078582</u>	<u>001</u>	Jan 15, 2009
<u>AA</u>		TARO	<u>100MG/ML</u>	<u>A078774</u>	<u>001</u>	Feb 10, 2009

TABLET; ORAL

KEPPRA

<u>AB</u>	<u>+</u>	UCB INC	<u>250MG</u>	<u>N021035</u>	<u>001</u>	Nov 30, 1999
<u>AB</u>	<u>+</u>		<u>500MG</u>	<u>N021035</u>	<u>002</u>	Nov 30, 1999
<u>AB</u>	<u>+</u>		<u>750MG</u>	<u>N021035</u>	<u>003</u>	Nov 30, 1999
<u>AB</u>	<u>+!</u>		<u>1GM</u>	<u>N021035</u>	<u>004</u>	Jan 06, 2006

LEVETIRACETAM

<u>AB</u>		ALKEM LABS LTD	<u>250MG</u>	<u>A216375</u>	<u>001</u>	May 27, 2022
<u>AB</u>			<u>500MG</u>	<u>A216375</u>	<u>002</u>	May 27, 2022
<u>AB</u>			<u>750MG</u>	<u>A216375</u>	<u>003</u>	May 27, 2022
<u>AB</u>			<u>1GM</u>	<u>A216375</u>	<u>004</u>	May 27, 2022
<u>AB</u>		AUROBINDO PHARMA	<u>250MG</u>	<u>A078993</u>	<u>001</u>	Jan 15, 2009
<u>AB</u>			<u>500MG</u>	<u>A078993</u>	<u>002</u>	Jan 15, 2009
<u>AB</u>			<u>750MG</u>	<u>A078993</u>	<u>003</u>	Jan 15, 2009
<u>AB</u>			<u>1GM</u>	<u>A078993</u>	<u>004</u>	Jan 15, 2009
<u>AB</u>		CHARTWELL RX	<u>250MG</u>	<u>A201293</u>	<u>001</u>	Jun 14, 2011
<u>AB</u>			<u>500MG</u>	<u>A201293</u>	<u>002</u>	Jun 14, 2011
<u>AB</u>			<u>750MG</u>	<u>A201293</u>	<u>003</u>	Jun 14, 2011
<u>AB</u>			<u>1GM</u>	<u>A201293</u>	<u>004</u>	Jun 14, 2011
<u>AB</u>		DR REDDYS LABS LTD	<u>250MG</u>	<u>A076920</u>	<u>001</u>	Jan 15, 2009
<u>AB</u>			<u>500MG</u>	<u>A076920</u>	<u>002</u>	Jan 15, 2009
<u>AB</u>			<u>750MG</u>	<u>A076920</u>	<u>003</u>	Jan 15, 2009
<u>AB</u>			<u>1GM</u>	<u>A078904</u>	<u>001</u>	Jan 15, 2009
<u>AB</u>		HETERO LABS LTD III	<u>250MG</u>	<u>A090515</u>	<u>001</u>	Oct 08, 2010
<u>AB</u>			<u>500MG</u>	<u>A090515</u>	<u>002</u>	Oct 08, 2010
<u>AB</u>			<u>750MG</u>	<u>A090515</u>	<u>003</u>	Oct 08, 2010
<u>AB</u>			<u>1GM</u>	<u>A090515</u>	<u>004</u>	Oct 08, 2010
<u>AB</u>		INGENUS PHARMS LLC	<u>250MG</u>	<u>A079042</u>	<u>001</u>	Jan 15, 2009
<u>AB</u>			<u>500MG</u>	<u>A079042</u>	<u>002</u>	Jan 15, 2009
<u>AB</u>			<u>750MG</u>	<u>A079042</u>	<u>003</u>	Jan 15, 2009
<u>AB</u>			<u>1GM</u>	<u>A079042</u>	<u>004</u>	Jan 15, 2009
<u>AB</u>		LUPIN	<u>250MG</u>	<u>A078154</u>	<u>001</u>	Jan 15, 2009
<u>AB</u>			<u>500MG</u>	<u>A078154</u>	<u>002</u>	Jan 15, 2009
<u>AB</u>			<u>750MG</u>	<u>A078154</u>	<u>003</u>	Jan 15, 2009
<u>AB</u>			<u>1GM</u>	<u>A090025</u>	<u>001</u>	Jan 15, 2009
<u>AB</u>		MSN	<u>250MG</u>	<u>A214815</u>	<u>001</u>	Oct 07, 2022
<u>AB</u>			<u>500MG</u>	<u>A214815</u>	<u>002</u>	Oct 07, 2022
<u>AB</u>			<u>750MG</u>	<u>A214815</u>	<u>003</u>	Oct 07, 2022
<u>AB</u>			<u>1GM</u>	<u>A214815</u>	<u>004</u>	Oct 07, 2022
<u>AB</u>		MYLAN	<u>500MG</u>	<u>A076919</u>	<u>002</u>	Nov 04, 2008
<u>AB</u>			<u>750MG</u>	<u>A076919</u>	<u>003</u>	Nov 04, 2008
<u>AB</u>			<u>1GM</u>	<u>A090261</u>	<u>001</u>	Dec 08, 2009
<u>AB</u>		ORBION PHARMS	<u>250MG</u>	<u>A078526</u>	<u>001</u>	Jan 15, 2009
<u>AB</u>			<u>500MG</u>	<u>A078526</u>	<u>002</u>	Jan 15, 2009
<u>AB</u>			<u>750MG</u>	<u>A078526</u>	<u>003</u>	Jan 15, 2009
<u>AB</u>			<u>1GM</u>	<u>A090484</u>	<u>001</u>	Aug 05, 2010
<u>AB</u>		OXFORD PHARMS	<u>250MG</u>	<u>A077319</u>	<u>001</u>	Mar 20, 2009
<u>AB</u>			<u>500MG</u>	<u>A077319</u>	<u>002</u>	Mar 20, 2009

PRESCRIPTION DRUG PRODUCT LIST

LEVETIRACETAM

TABLET; ORAL

LEVETIRACETAM

<u>AB</u>		<u>750MG</u>	<u>A077319</u>	<u>003</u>	Mar 20, 2009
<u>AB</u>	PRINSTON INC	<u>250MG</u>	<u>A078106</u>	<u>001</u>	Feb 10, 2009
<u>AB</u>		<u>500MG</u>	<u>A078106</u>	<u>002</u>	Feb 10, 2009
<u>AB</u>		<u>750MG</u>	<u>A078106</u>	<u>003</u>	Feb 10, 2009
<u>AB</u>		<u>1GM</u>	<u>A078106</u>	<u>004</u>	Feb 10, 2009
<u>AB</u>	RISING	<u>250MG</u>	<u>A090767</u>	<u>001</u>	Jul 28, 2010
<u>AB</u>		<u>500MG</u>	<u>A090767</u>	<u>002</u>	Jul 28, 2010
<u>AB</u>		<u>750MG</u>	<u>A090767</u>	<u>003</u>	Jul 28, 2010
<u>AB</u>		<u>1GM</u>	<u>A090767</u>	<u>004</u>	Jul 28, 2010
<u>AB</u>	SCIEGEN PHARMS INC	<u>250MG</u>	<u>A215069</u>	<u>003</u>	May 27, 2022
<u>AB</u>		<u>500MG</u>	<u>A215069</u>	<u>004</u>	May 27, 2022
<u>AB</u>		<u>750MG</u>	<u>A215069</u>	<u>001</u>	Jun 11, 2021
<u>AB</u>		<u>1GM</u>	<u>A215069</u>	<u>002</u>	Jun 11, 2021
<u>AB</u>	SECAN PHARMS	<u>250MG</u>	<u>A205102</u>	<u>001</u>	Dec 16, 2015
<u>AB</u>		<u>750MG</u>	<u>A205102</u>	<u>002</u>	Dec 16, 2015
<u>AB</u>	STRIDES PHARMA	<u>250MG</u>	<u>A078042</u>	<u>001</u>	Jan 15, 2009
<u>AB</u>		<u>500MG</u>	<u>A078042</u>	<u>002</u>	Jan 15, 2009
<u>AB</u>		<u>750MG</u>	<u>A078042</u>	<u>003</u>	Jan 15, 2009
<u>AB</u>		<u>1GM</u>	<u>A078042</u>	<u>004</u>	Jan 15, 2009
<u>AB</u>	TARO	<u>250MG</u>	<u>A078960</u>	<u>004</u>	Feb 01, 2010
<u>AB</u>		<u>500MG</u>	<u>A078960</u>	<u>003</u>	Feb 01, 2010
<u>AB</u>		<u>750MG</u>	<u>A078960</u>	<u>002</u>	Feb 01, 2010
<u>AB</u>		<u>1GM</u>	<u>A078960</u>	<u>001</u>	Feb 01, 2010
<u>AB</u>	TORRENT PHARMS	<u>250MG</u>	<u>A078858</u>	<u>001</u>	Jan 15, 2009
<u>AB</u>		<u>500MG</u>	<u>A078858</u>	<u>002</u>	Jan 15, 2009
<u>AB</u>		<u>750MG</u>	<u>A078858</u>	<u>003</u>	Jan 15, 2009
<u>AB</u>		<u>1GM</u>	<u>A078858</u>	<u>004</u>	Jan 15, 2009
<u>AB</u>	VIWIT PHARM	<u>250MG</u>	<u>A078869</u>	<u>001</u>	Mar 13, 2009
<u>AB</u>		<u>500MG</u>	<u>A078869</u>	<u>002</u>	Mar 13, 2009
<u>AB</u>		<u>750MG</u>	<u>A078869</u>	<u>003</u>	Mar 13, 2009
<u>AB</u>		<u>1GM</u>	<u>A078869</u>	<u>004</u>	Mar 13, 2009
<u>AB</u>	ZHEJIANG JINGXIN	<u>250MG</u>	<u>A091491</u>	<u>001</u>	Dec 14, 2010
<u>AB</u>		<u>500MG</u>	<u>A091491</u>	<u>002</u>	Dec 14, 2010
<u>AB</u>		<u>750MG</u>	<u>A091491</u>	<u>003</u>	Dec 14, 2010
<u>AB</u>		<u>1GM</u>	<u>A091491</u>	<u>004</u>	Dec 14, 2010
<u>AB</u>	ZYDUS PHARMS USA INC	<u>250MG</u>	<u>A078918</u>	<u>001</u>	Apr 29, 2009
<u>AB</u>		<u>1GM</u>	<u>A078918</u>	<u>002</u>	Apr 29, 2009

TABLET, EXTENDED RELEASE; ORAL

KEPPRA XR

<u>AB</u>	+ UCB INC	<u>500MG</u>	<u>N022285</u>	<u>001</u>	Sep 12, 2008
<u>AB</u>	+!	<u>750MG</u>	<u>N022285</u>	<u>002</u>	Feb 12, 2009

LEVETIRACETAM

<u>AB</u>	ACTAVIS LABS FL INC	<u>500MG</u>	<u>A091093</u>	<u>001</u>	Sep 12, 2011
<u>AB</u>		<u>750MG</u>	<u>A091093</u>	<u>002</u>	Sep 12, 2011
<u>AB</u>	AIPING PHARM INC	<u>500MG</u>	<u>A204754</u>	<u>001</u>	Aug 26, 2016
<u>AB</u>		<u>750MG</u>	<u>A204754</u>	<u>002</u>	Aug 26, 2016
<u>AB</u>	ANDA REPOSITORY	<u>500MG</u>	<u>A204511</u>	<u>001</u>	Feb 23, 2016
<u>AB</u>		<u>750MG</u>	<u>A204511</u>	<u>002</u>	Feb 23, 2016
<u>AB</u>	APOTEX INC	<u>500MG</u>	<u>A091261</u>	<u>001</u>	Sep 12, 2011
<u>AB</u>		<u>750MG</u>	<u>A091261</u>	<u>002</u>	Sep 12, 2011
<u>AB</u>	HISUN PHARM HANGZHOU	<u>500MG</u>	<u>A207175</u>	<u>001</u>	Sep 28, 2017
<u>AB</u>		<u>750MG</u>	<u>A207175</u>	<u>002</u>	Sep 28, 2017
<u>AB</u>	LUPIN LTD	<u>500MG</u>	<u>A091399</u>	<u>001</u>	Sep 12, 2011
<u>AB</u>		<u>750MG</u>	<u>A091399</u>	<u>002</u>	Sep 12, 2011
<u>AB</u>	OVERSEAS	<u>500MG</u>	<u>A212688</u>	<u>002</u>	May 05, 2023
<u>AB</u>		<u>750MG</u>	<u>A212688</u>	<u>001</u>	Jun 11, 2020
<u>AB</u>	PRINSTON INC	<u>500MG</u>	<u>A203468</u>	<u>001</u>	May 21, 2015
<u>AB</u>		<u>750MG</u>	<u>A203468</u>	<u>002</u>	May 21, 2015
<u>AB</u>	SCIEGEN PHARMS INC	<u>500MG</u>	<u>A205130</u>	<u>001</u>	Nov 27, 2020
<u>AB</u>		<u>750MG</u>	<u>A205130</u>	<u>002</u>	Nov 27, 2020
<u>AB</u>	APOTEX	1GM	A202958	001	Feb 25, 2015

TABLET, FOR SUSPENSION; ORAL

SPRITAM

+	APRECIA PHARMS	250MG	N207958	001	Jul 31, 2015
+		500MG	N207958	002	Jul 31, 2015
+		750MG	N207958	003	Jul 31, 2015
+		1GM	N207958	004	Jul 31, 2015

PRESCRIPTION DRUG PRODUCT LIST

LEVOBUNOLOL HYDROCHLORIDE

SOLUTION/DROPS;OPHTHALMIC

BETAGAN

AT	+ !	ALLERGAN	0.5%	N019219	002	Dec 19, 1985
-----------	------------	----------	-------------	----------------	------------	--------------

LEVOBUNOLOL HYDROCHLORIDE

AT		BAUSCH AND LOMB	0.5%	A074326	001	Mar 04, 1994
-----------	--	-----------------	-------------	----------------	------------	--------------

LEVOCARNITINE

INJECTABLE; INJECTION

CARNITOR

AP	+ !	LEADIANT BIOSCI INC	200MG/ML	N020182	001	Dec 16, 1992
-----------	------------	---------------------	-----------------	----------------	------------	--------------

LEVOCARNITINE

AP		AM REGENT	200MG/ML	A075861	001	Jun 22, 2001
-----------	--	-----------	-----------------	----------------	------------	--------------

AP		HIKMA	200MG/ML	A075567	001	Mar 29, 2001
-----------	--	-------	-----------------	----------------	------------	--------------

SOLUTION; ORAL

CARNITOR

AA	+ !	LEADIANT BIOSCI INC	1GM/10ML	N019257	001	Apr 10, 1986
-----------	------------	---------------------	-----------------	----------------	------------	--------------

CARNITOR SF

AA	+	LEADIANT BIOSCI INC	1GM/10ML	N019257	002	Mar 28, 2007
-----------	----------	---------------------	-----------------	----------------	------------	--------------

LEVOCARNITINE

AA		NOVITIUM PHARMA	1GM/10ML	A211676	001	Aug 14, 2019
-----------	--	-----------------	-----------------	----------------	------------	--------------

AA		RISING	1GM/10ML	A076851	001	Aug 10, 2004
-----------	--	--------	-----------------	----------------	------------	--------------

AA		SCIEGEN PHARMS INC	1GM/10ML	A212533	001	Nov 10, 2021
-----------	--	--------------------	-----------------	----------------	------------	--------------

LEVOCARNITINE SF

AA		NOVITIUM PHARMA	1GM/10ML	A211676	002	Aug 14, 2019
-----------	--	-----------------	-----------------	----------------	------------	--------------

TABLET; ORAL

CARNITOR

AB	+ !	LEADIANT BIOSCI INC	330MG	N018948	001	Dec 27, 1985
-----------	------------	---------------------	--------------	----------------	------------	--------------

LEVOCARNITINE

AB		NOVITIUM PHARMA	330MG	A216384	001	Dec 09, 2022
-----------	--	-----------------	--------------	----------------	------------	--------------

AB		RISING	330MG	A076858	001	Sep 20, 2004
-----------	--	--------	--------------	----------------	------------	--------------

LEVOCETIRIZINE DIHYDROCHLORIDE

SOLUTION; ORAL

LEVOCETIRIZINE DIHYDROCHLORIDE

AA		CHARTWELL MOLECULAR	2.5MG/5ML	A204599	001	May 15, 2017
-----------	--	---------------------	------------------	----------------	------------	--------------

AA	!	PADAGIS US	2.5MG/5ML	A091263	001	Nov 07, 2011
-----------	----------	------------	------------------	----------------	------------	--------------

AA		TARO	2.5MG/5ML	A202673	001	Jul 26, 2013
-----------	--	------	------------------	----------------	------------	--------------

LEVOCETIRIZINE HYDROCHLORIDE

AA		HETERO LABS LTD III	2.5MG/5ML	A210914	001	Apr 01, 2019
-----------	--	---------------------	------------------	----------------	------------	--------------

TABLET; ORAL

LEVOCETIRIZINE DIHYDROCHLORIDE

AB		DR REDDYS LABS LTD	5MG	A090392	001	Feb 24, 2011
-----------	--	--------------------	------------	----------------	------------	--------------

AB	!	GLENMARK PHARMS LTD	5MG	A090385	001	Feb 24, 2011
-----------	----------	---------------------	------------	----------------	------------	--------------

AB		HETERO LABS LTD III	5MG	A091264	001	Jun 29, 2012
-----------	--	---------------------	------------	----------------	------------	--------------

AB		MACLEODS PHARMS LTD	5MG	A205564	001	Jan 11, 2016
-----------	--	---------------------	------------	----------------	------------	--------------

AB		MICRO LABS LTD	5MG	A202046	001	Sep 17, 2013
-----------	--	----------------	------------	----------------	------------	--------------

INDIA

AB		SCIEGEN PHARMS INC	5MG	A203646	001	Sep 09, 2014
-----------	--	--------------------	------------	----------------	------------	--------------

AB		SUN PHARM	5MG	A090362	001	Jan 31, 2013
-----------	--	-----------	------------	----------------	------------	--------------

AB		SYNTHON PHARMS	5MG	A090229	001	Nov 26, 2010
-----------	--	----------------	------------	----------------	------------	--------------

AB		TEVA PHARMS	5MG	A090199	001	Aug 22, 2011
-----------	--	-------------	------------	----------------	------------	--------------

LEVODOPA

POWDER; INHALATION

INBRIJA

+ !		MERZ PHARMS	42MG	N209184	001	Dec 21, 2018
------------	--	-------------	-------------	----------------	------------	--------------

LEVOFLOXACIN

INJECTABLE; INJECTION

LEVOFLOXACIN

AP		GLAND PHARMA LTD	EQ 500MG/20ML (EQ 25MG/ML)	A205540	001	Apr 22, 2020
-----------	--	------------------	-----------------------------------	----------------	------------	--------------

AP			EQ 750MG/30ML (EQ 25MG/ML)	A205540	002	Apr 22, 2020
-----------	--	--	-----------------------------------	----------------	------------	--------------

AP	!	RISING	EQ 500MG/20ML (EQ 25MG/ML)	A091644	001	Jun 20, 2011
-----------	----------	--------	-----------------------------------	----------------	------------	--------------

AP	!		EQ 750MG/30ML (EQ 25MG/ML)	A091644	002	Jun 20, 2011
-----------	----------	--	-----------------------------------	----------------	------------	--------------

LEVOFLOXACIN IN DEXTROSE 5% IN PLASTIC CONTAINER

AP		FRESENIUS KABI USA	EQ 250MG/50ML (EQ 5MG/ML)	A200674	001	Jun 19, 2013
-----------	--	--------------------	----------------------------------	----------------	------------	--------------

AP			EQ 500MG/100ML (EQ 5MG/ML)	A200674	002	Jun 19, 2013
-----------	--	--	-----------------------------------	----------------	------------	--------------

AP			EQ 750MG/150ML (EQ 5MG/ML)	A200674	003	Jun 19, 2013
-----------	--	--	-----------------------------------	----------------	------------	--------------

AP		GLAND PHARMA LTD	EQ 250MG/50ML (EQ 5MG/ML)	A206908	001	Dec 30, 2020
-----------	--	------------------	----------------------------------	----------------	------------	--------------

AP			EQ 500MG/100ML (EQ 5MG/ML)	A206908	002	Dec 30, 2020
-----------	--	--	-----------------------------------	----------------	------------	--------------

AP			EQ 750MG/150ML (EQ 5MG/ML)	A206908	003	Dec 30, 2020
-----------	--	--	-----------------------------------	----------------	------------	--------------

AP		HIKMA FARMACEUTICA	EQ 250MG/50ML (EQ 5MG/ML)	A091375	001	Sep 16, 2011
-----------	--	--------------------	----------------------------------	----------------	------------	--------------

AP			EQ 500MG/100ML (EQ 5MG/ML)	A091375	002	Sep 16, 2011
-----------	--	--	-----------------------------------	----------------	------------	--------------

AP			EQ 750MG/150ML (EQ 5MG/ML)	A091375	003	Sep 16, 2011
-----------	--	--	-----------------------------------	----------------	------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

LEVOFLOXACIN

INJECTABLE; INJECTION

LEVOFLOXACIN IN DEXTROSE 5% IN PLASTIC CONTAINER

<u>AP</u>	HOSPIRA	<u>EQ 250MG/50ML (EQ 5MG/ML)</u>	<u>A078579 001</u>	Sep 03, 2015
<u>AP</u>		<u>EQ 500MG/100ML (EQ 5MG/ML)</u>	<u>A078579 002</u>	Sep 03, 2015
<u>AP</u>		<u>EQ 750MG/150ML (EQ 5MG/ML)</u>	<u>A078579 003</u>	Sep 03, 2015
<u>AP</u>	! INFORLIFE	<u>EQ 250MG/50ML (EQ 5MG/ML)</u>	<u>A090343 001</u>	Jul 07, 2011
<u>AP</u>	!	<u>EQ 500MG/100ML (EQ 5MG/ML)</u>	<u>A090343 002</u>	Jul 07, 2011
<u>AP</u>	!	<u>EQ 750MG/150ML (EQ 5MG/ML)</u>	<u>A090343 003</u>	Jul 07, 2011

SOLUTION; ORAL

LEVOFLOXACIN

<u>AA</u>	LANNETT CO INC	<u>250MG/10ML</u>	<u>A205222 001</u>	May 25, 2018
<u>AA</u>	! NOVITIUM PHARMA	<u>250MG/10ML</u>	<u>A091678 001</u>	Jun 20, 2011

SOLUTION/DROPS; OPHTHALMIC

LEVOFLOXACIN

<u>AT</u>	! RISING	<u>0.5%</u>	<u>A077700 001</u>	Dec 20, 2010
<u>AT</u>	RUBICON	<u>0.5%</u>	<u>A078282 001</u>	Dec 20, 2010
	MICRO LABS LTD INDIA	1.5%	A205600 001	Feb 27, 2019

TABLET; ORAL

LEVOFLOXACIN

<u>AB</u>	APOTEX INC	<u>250MG</u>	<u>A090787 001</u>	Sep 29, 2011
<u>AB</u>		<u>500MG</u>	<u>A090787 002</u>	Sep 29, 2011
<u>AB</u>		<u>750MG</u>	<u>A090787 003</u>	Sep 29, 2011
<u>AB</u>	AUROBINDO PHARMA LTD	<u>250MG</u>	<u>A201043 001</u>	Jun 20, 2011
<u>AB</u>		<u>500MG</u>	<u>A201043 002</u>	Jun 20, 2011
<u>AB</u>	!	<u>750MG</u>	<u>A201043 003</u>	Jun 20, 2011
<u>AB</u>	CHARTWELL MOLECULAR	<u>250MG</u>	<u>A076890 001</u>	Mar 30, 2012
<u>AB</u>		<u>500MG</u>	<u>A076890 002</u>	Mar 30, 2012
<u>AB</u>		<u>750MG</u>	<u>A076890 003</u>	Mar 30, 2012
<u>AB</u>	DR REDDYS LABS INC	<u>250MG</u>	<u>A076710 001</u>	Jun 20, 2011
<u>AB</u>		<u>500MG</u>	<u>A076710 002</u>	Jun 20, 2011
<u>AB</u>		<u>750MG</u>	<u>A076710 003</u>	Jun 20, 2011
<u>AB</u>	GLENMARK PHARMS LTD	<u>250MG</u>	<u>A200250 001</u>	Jun 20, 2011
<u>AB</u>		<u>500MG</u>	<u>A200250 002</u>	Jun 20, 2011
<u>AB</u>		<u>750MG</u>	<u>A200250 003</u>	Jun 20, 2011
<u>AB</u>	HEC PHARM	<u>250MG</u>	<u>A204968 001</u>	Feb 05, 2019
<u>AB</u>		<u>500MG</u>	<u>A204968 002</u>	Feb 05, 2019
<u>AB</u>		<u>750MG</u>	<u>A204968 003</u>	Feb 05, 2019
<u>AB</u>	HETERO LABS LTD V	<u>250MG</u>	<u>A202801 001</u>	Jan 08, 2015
<u>AB</u>		<u>500MG</u>	<u>A202801 002</u>	Jan 08, 2015
<u>AB</u>		<u>750MG</u>	<u>A202801 003</u>	Jan 08, 2015
<u>AB</u>	LUPIN	<u>250MG</u>	<u>A078424 001</u>	Jun 20, 2011
<u>AB</u>		<u>500MG</u>	<u>A078424 002</u>	Jun 20, 2011
<u>AB</u>		<u>750MG</u>	<u>A078424 003</u>	Jun 20, 2011
<u>AB</u>	MACLEODS PHARMS LTD	<u>250MG</u>	<u>A200839 001</u>	Mar 22, 2012
<u>AB</u>		<u>500MG</u>	<u>A200839 002</u>	Mar 22, 2012
<u>AB</u>		<u>750MG</u>	<u>A200839 003</u>	Mar 22, 2012
<u>AB</u>	ORBION PHARMS	<u>250MG</u>	<u>A202200 001</u>	Jan 30, 2012
<u>AB</u>		<u>500MG</u>	<u>A202200 002</u>	Jan 30, 2012
<u>AB</u>		<u>750MG</u>	<u>A202200 003</u>	Jan 30, 2012
<u>AB</u>	TEVA	<u>250MG</u>	<u>A076361 001</u>	Jun 20, 2011
<u>AB</u>		<u>500MG</u>	<u>A076361 002</u>	Jun 20, 2011
<u>AB</u>		<u>750MG</u>	<u>A076361 003</u>	Jun 20, 2011
<u>AB</u>	ZYDUS LIFESCIENCES	<u>250MG</u>	<u>A077652 001</u>	Sep 07, 2012
<u>AB</u>		<u>500MG</u>	<u>A077652 002</u>	Sep 07, 2012
<u>AB</u>		<u>750MG</u>	<u>A077652 003</u>	Sep 07, 2012

LEVOKETOCONAZOLE

TABLET; ORAL

RECORLEV

+	!	STRONGBRIDGE	150MG	N214133 001	Dec 30, 2021
---	---	--------------	-------	-------------	--------------

LEVOLEUCOVORIN

POWDER; INTRAVENOUS

KHAPZORY

+	!	ACROTECH BIOPHARMA	175MG/VIAL	N211226 001	Oct 19, 2018
---	---	--------------------	------------	-------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

LEVOLEUCOVORIN CALCIUM

SOLUTION; INTRAVENOUS

LEVOLEUCOVORIN CALCIUM

<u>AP</u>	AMNEAL	<u>EQ 175MG BASE/17.5ML (EQ 10MG BASE/ML)</u>	<u>A207548 001</u>	Sep 08, 2017
<u>AP</u>	GLAND PHARMA LTD	<u>EQ 175MG BASE/17.5ML (EQ 10MG BASE/ML)</u>	<u>A210892 001</u>	Sep 14, 2018
<u>AP</u>	!	<u>EQ 250MG BASE/25ML (EQ 10MG BASE/ML)</u>	<u>A210892 002</u>	Sep 14, 2018
<u>AP</u>	HAINAN POLY PHARM	<u>EQ 175MG BASE/17.5ML (EQ 10MG BASE/ML)</u>	<u>A217314 001</u>	May 22, 2023
<u>AP</u>		<u>EQ 250MG BASE/25ML (EQ 10MG BASE/ML)</u>	<u>A217314 002</u>	May 22, 2023
<u>AP</u>	!	<u>EQ 175MG BASE/17.5ML (EQ 10MG BASE/ML)</u>	<u>A211002 001</u>	Aug 16, 2019
<u>AP</u>	SANDOZ	<u>EQ 175MG BASE/17.5ML (EQ 10MG BASE/ML)</u>	<u>A203563 001</u>	Mar 09, 2015

LEVOMILNACIPRAN HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

LEVOMILNACIPRAN HYDROCHLORIDE

<u>AB</u>	PRINSTON INC	<u>EQ 20MG BASE</u>	<u>A210771 001</u>	Mar 20, 2023
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A210771 002</u>	Mar 20, 2023
<u>AB</u>		<u>EQ 80MG BASE</u>	<u>A210771 003</u>	Mar 20, 2023
<u>AB</u>		<u>EQ 120MG BASE</u>	<u>A210771 004</u>	Mar 20, 2023
	FETZIMA			
	+ ABBVIE	EQ 20MG BASE	N204168 001	Jul 25, 2013
	+	EQ 40MG BASE	N204168 002	Jul 25, 2013
	+	EQ 80MG BASE	N204168 003	Jul 25, 2013
	+	EQ 120MG BASE	N204168 004	Jul 25, 2013

LEVONORGESTREL

SYSTEM; INTRAUTERINE

KYLEENA

+! BAYER HLTHCARE 19.5MG N208224 001 Sep 16, 2016

LILETTA

+! MEDICINES360 52MG N206229 001 Feb 26, 2015

MIRENA

+! BAYER HLTHCARE 52MG N021225 001 Dec 06, 2000

SKYLA

+! BAYER HLTHCARE 13.5MG N203159 001 Jan 09, 2013

LEVORPHANOL TARTRATE

TABLET; ORAL

LEVORPHANOL TARTRATE

<u>AB</u>	HIKMA	<u>2MG</u>	<u>A074278 001</u>	Mar 31, 2000
<u>AB</u>	NOVITIUM PHARMA	<u>2MG</u>	<u>A213479 001</u>	Jul 01, 2020
<u>AB</u>	!	<u>3MG</u>	<u>A213479 002</u>	Jan 12, 2021
<u>AB</u>	SPECGX LLC	<u>2MG</u>	<u>A212024 001</u>	Dec 13, 2019
<u>AB</u>	SUN PHARM INDS INC	<u>2MG</u>	<u>A213906 001</u>	Jun 17, 2021
<u>AB</u>		<u>3MG</u>	<u>A213906 002</u>	Apr 25, 2023
<u>AB</u>	VIRTUS	<u>2MG</u>	<u>A211484 001</u>	Dec 13, 2018
	NOVITIUM PHARMA	1MG	A213479 003	Jul 20, 2021

LEVOTHYROXINE SODIUM

CAPSULE; ORAL

TIROSINT

+ IBSA 0.013MG N021924 013 Aug 01, 2007

+ 0.025MG N021924 002 Oct 13, 2006

+ 0.0375MG N021924 014 Jun 22, 2022

+ 0.044MG N021924 015 Jun 22, 2022

+ 0.05MG N021924 003 Oct 13, 2006

+ 0.0625MG N021924 016 Jun 22, 2022

+ 0.075MG N021924 004 Oct 13, 2006

+ 0.088MG N021924 010 Oct 02, 2009

+ 0.1MG N021924 005 Oct 13, 2006

+ 0.112MG N021924 008 Oct 02, 2009

+ 0.125MG N021924 006 Oct 13, 2006

+ 0.137MG N021924 009 Oct 02, 2009

+ 0.15MG N021924 007 Oct 13, 2006

+ 0.175MG N021924 011 Apr 25, 2017

+! 0.2MG N021924 012 Apr 25, 2017

POWDER; INTRAVENOUS

LEVOTHYROXINE SODIUM

<u>AP</u>	+!	FRESENIUS KABI USA	<u>100MCG/VIAL</u>	<u>N202231 001</u>	Jun 24, 2011
<u>AP</u>	+!		<u>200MCG/VIAL</u>	<u>N202231 002</u>	Jun 24, 2011
<u>AP</u>	+!		<u>500MCG/VIAL</u>	<u>N202231 003</u>	Jun 24, 2011
<u>AP</u>		MAIA PHARMS INC	<u>100MCG/VIAL</u>	<u>A208749 001</u>	Dec 21, 2018
<u>AP</u>			<u>200MCG/VIAL</u>	<u>A208749 002</u>	Dec 21, 2018
<u>AP</u>			<u>500MCG/VIAL</u>	<u>A208749 003</u>	Dec 21, 2018
<u>AP</u>		PIRAMAL CRITICAL	<u>100MCG/VIAL</u>	<u>A206163 001</u>	Jun 29, 2016

PRESCRIPTION DRUG PRODUCT LIST

LEVOTHYROXINE SODIUM

POWDER; INTRAVENOUS

LEVOTHYROXINE SODIUM

AP		<u>500MCG/VIAL</u>	<u>A206163 002</u>	Jun 29, 2016
AP	XIROMED	<u>100MCG/VIAL</u>	<u>A217495 001</u>	Nov 29, 2024
AP	ZYDUS PHARMS	<u>100MCG/VIAL</u>	<u>A217066 001</u>	Mar 24, 2023
AP		<u>200MCG/VIAL</u>	<u>A217066 002</u>	Mar 24, 2023
AP		<u>500MCG/VIAL</u>	<u>A217066 003</u>	Mar 24, 2023

SOLUTION; INTRAVENOUS

LEVOTHYROXINE SODIUM

+!	FRESENIUS KABI USA	100MCG/5ML (20MCG/ML)	N210632 001	Apr 11, 2019
+!		200MCG/5ML (40MCG/ML)	N210632 002	Apr 11, 2019
+!		500MCG/5ML (100MCG/ML)	N210632 003	Apr 11, 2019
+!	HIKMA	100MCG/ML	N214253 001	May 17, 2021

SOLUTION; ORAL

ERMEZA

+!	MYLAN	150MCG/5ML	N215809 001	Apr 29, 2022
----	-------	------------	-------------	--------------

THYQUIDITY

+!	AZURITY	100MCG/5ML	N214047 001	Nov 30, 2020
----	---------	------------	-------------	--------------

TIROSINT-SOL

+	IBSA	13MCG/ML	N206977 001	Dec 15, 2016
+		25MCG/ML	N206977 002	Dec 15, 2016
+		37.5MCG/ML	N206977 013	Jan 13, 2021
+		44MCG/ML	N206977 014	Jan 13, 2021
+		50MCG/ML	N206977 003	Dec 15, 2016
+		62.5MCG/ML	N206977 015	Jan 13, 2021
+		75MCG/ML	N206977 004	Dec 15, 2016
+		88MCG/ML	N206977 005	Dec 15, 2016
+		100MCG/ML	N206977 006	Dec 15, 2016
+		112MCG/ML	N206977 007	Dec 15, 2016
+		125MCG/ML	N206977 008	Dec 15, 2016
+		137MCG/ML	N206977 009	Dec 15, 2016
+		150MCG/ML	N206977 010	Dec 15, 2016
+		175MCG/ML	N206977 011	Dec 15, 2016
+!		200MCG/ML	N206977 012	Dec 15, 2016

LEVOTHYROXINE SODIUM **

**See current Annual Edition, 1.8 Description of Special Situations, Levothyroxine Sodium

TABLET; ORAL

LEVOTHYROXINE SODIUM

-->	MACLEODS PHARMS LTD	--> <u>AB1, AB2</u>	<u>0.025MG</u>	A211417 001	Dec 21, 2022
-->		--> <u>AB1, AB2</u>	<u>0.05MG</u>	A211417 002	Dec 21, 2022
-->		--> <u>AB1, AB2</u>	<u>0.075MG</u>	A211417 003	Dec 21, 2022
-->		--> <u>AB1, AB2</u>	<u>0.088MG</u>	A211417 004	Dec 21, 2022
-->		--> <u>AB1, AB2</u>	<u>0.1MG</u>	A211417 005	Dec 21, 2022
-->		--> <u>AB1, AB2</u>	<u>0.112MG</u>	A211417 006	Dec 21, 2022
-->		--> <u>AB1, AB2</u>	<u>0.125MG</u>	A211417 007	Dec 21, 2022
-->		--> <u>AB1, AB2</u>	<u>0.137MG</u>	A211417 008	Dec 21, 2022
-->		--> <u>AB1, AB2</u>	<u>0.15MG</u>	A211417 009	Dec 21, 2022
-->		--> <u>AB1, AB2</u>	<u>0.175MG</u>	A211417 010	Dec 21, 2022
-->		--> <u>AB1, AB2</u>	<u>0.2MG</u>	A211417 011	Dec 21, 2022
-->		--> <u>AB1, AB2</u>	<u>0.3MG</u>	A211417 012	Dec 21, 2022

SYNTHROID

-->	+	ABEVIE	--> <u>AB1, AB2</u>	<u>0.025MG</u>	N021402 001	Jul 24, 2002
-->	+		--> <u>AB1, AB2</u>	<u>0.05MG</u>	N021402 002	Jul 24, 2002
-->	+		--> <u>AB1, AB2</u>	<u>0.075MG</u>	N021402 003	Jul 24, 2002
-->	+		--> <u>AB1, AB2</u>	<u>0.088MG</u>	N021402 004	Jul 24, 2002
-->	+		--> <u>AB1, AB2</u>	<u>0.1MG</u>	N021402 005	Jul 24, 2002
-->	+		--> <u>AB1, AB2</u>	<u>0.112MG</u>	N021402 006	Jul 24, 2002
-->	+		--> <u>AB1, AB2</u>	<u>0.125MG</u>	N021402 007	Jul 24, 2002
-->	+		--> <u>AB1, AB2</u>	<u>0.137MG</u>	N021402 008	Jul 24, 2002
-->	+		--> <u>AB1, AB2</u>	<u>0.15MG</u>	N021402 009	Jul 24, 2002
-->	+		--> <u>AB1, AB2</u>	<u>0.175MG</u>	N021402 010	Jul 24, 2002
-->	+		--> <u>AB1, AB2</u>	<u>0.2MG</u>	N021402 012	Jul 24, 2002
-->	+!		--> <u>AB1, AB2</u>	<u>0.3MG</u>	N021402 011	Jul 24, 2002

LEVO-T

-->	CEDIPROF INC	--> <u>AB1, AB2, AB3</u>	<u>0.025MG</u>	N021342 001	Mar 01, 2002
-->		--> <u>AB1, AB2, AB3</u>	<u>0.05MG</u>	N021342 002	Mar 01, 2002
-->		--> <u>AB1, AB2, AB3</u>	<u>0.075MG</u>	N021342 003	Mar 01, 2002
-->		--> <u>AB1, AB2, AB3</u>	<u>0.088MG</u>	N021342 004	Mar 01, 2002

PRESCRIPTION DRUG PRODUCT LIST

LEVOTHYROXINE SODIUM **

**See current Annual Edition, 1.8 Description of Special Situations, Levothyroxine Sodium

TABLET;ORAL

LEVO-T

-->	--> <u>AB1,AB2,AB3</u>	<u>0.1MG</u>	N021342 005	Mar 01, 2002
-->	--> <u>AB1,AB2,AB3</u>	<u>0.112MG</u>	N021342 006	Mar 01, 2002
-->	--> <u>AB1,AB2,AB3</u>	<u>0.125MG</u>	N021342 007	Mar 01, 2002
-->	--> <u>AB1,AB2,AB3</u>	<u>0.137MG</u>	N021342 012	Dec 08, 2003
-->	--> <u>AB1,AB2,AB3</u>	<u>0.15MG</u>	N021342 008	Mar 01, 2002
-->	--> <u>AB1,AB2,AB3</u>	<u>0.175MG</u>	N021342 009	Mar 01, 2002
-->	--> <u>AB1,AB2,AB3</u>	<u>0.2MG</u>	N021342 010	Mar 01, 2002
-->	--> <u>AB1,AB2,AB3</u>	<u>0.3MG</u>	N021342 011	Mar 01, 2002

LEVOTHYROXINE SODIUM

-->	LUPIN	--> <u>AB1,AB2,AB3</u>	<u>0.025MG</u>	A209713 001	Jan 18, 2019
-->		--> <u>AB1,AB2,AB3</u>	<u>0.05MG</u>	A209713 002	Jan 18, 2019
-->		--> <u>AB1,AB2,AB3</u>	<u>0.075MG</u>	A209713 003	Jan 18, 2019
-->		--> <u>AB1,AB2,AB3</u>	<u>0.088MG</u>	A209713 004	Jan 18, 2019
-->		--> <u>AB1,AB2,AB3</u>	<u>0.1MG</u>	A209713 005	Jan 18, 2019
-->		--> <u>AB1,AB2,AB3</u>	<u>0.112MG</u>	A209713 006	Jan 18, 2019
-->		--> <u>AB1,AB2,AB3</u>	<u>0.125MG</u>	A209713 007	Jan 18, 2019
-->		--> <u>AB1,AB2,AB3</u>	<u>0.137MG</u>	A209713 008	Jan 18, 2019
-->		--> <u>AB1,AB2,AB3</u>	<u>0.15MG</u>	A209713 009	Jan 18, 2019
-->		--> <u>AB1,AB2,AB3</u>	<u>0.175MG</u>	A209713 010	Jan 18, 2019
-->		--> <u>AB1,AB2,AB3</u>	<u>0.2MG</u>	A209713 011	Jan 18, 2019
-->		--> <u>AB1,AB2,AB3</u>	<u>0.3MG</u>	A209713 012	Jan 18, 2019

UNITHROID

-->	+ STEVENS J	--> <u>AB1,AB2,AB3</u>	<u>0.025MG</u>	N021210 001	Aug 21, 2000
-->	+	--> <u>AB1,AB2,AB3</u>	<u>0.05MG</u>	N021210 002	Aug 21, 2000
-->	+	--> <u>AB1,AB2,AB3</u>	<u>0.075MG</u>	N021210 003	Aug 21, 2000
-->	+	--> <u>AB1,AB2,AB3</u>	<u>0.088MG</u>	N021210 004	Aug 21, 2000
-->	+	--> <u>AB1,AB2,AB3</u>	<u>0.1MG</u>	N021210 005	Aug 21, 2000
-->	+	--> <u>AB1,AB2,AB3</u>	<u>0.112MG</u>	N021210 006	Aug 21, 2000
-->	+	--> <u>AB1,AB2,AB3</u>	<u>0.125MG</u>	N021210 007	Aug 21, 2000
-->	+	--> <u>AB1,AB2,AB3</u>	<u>0.137MG</u>	N021210 012	Feb 08, 2008
-->	+	--> <u>AB1,AB2,AB3</u>	<u>0.15MG</u>	N021210 008	Aug 21, 2000
-->	+	--> <u>AB1,AB2,AB3</u>	<u>0.175MG</u>	N021210 009	Aug 21, 2000
-->	+	--> <u>AB1,AB2,AB3</u>	<u>0.2MG</u>	N021210 010	Aug 21, 2000
-->	+!	--> <u>AB1,AB2,AB3</u>	<u>0.3MG</u>	N021210 011	Aug 21, 2000

LEVOLET

-->	GENUS LIFESCIENCES	-->	<u>0.025MG</u>	N021137 001	Jun 06, 2003
-->		<u>AB1,AB2,AB3,AB4</u>			
-->		-->	<u>0.05MG</u>	N021137 002	Jun 06, 2003
-->		<u>AB1,AB2,AB3,AB4</u>			
-->		-->	<u>0.075MG</u>	N021137 003	Jun 06, 2003
-->		<u>AB1,AB2,AB3,AB4</u>			
-->		-->	<u>0.088MG</u>	N021137 004	Jun 06, 2003
-->		<u>AB1,AB2,AB3,AB4</u>			
-->		--> <u>AB1,AB2,</u>	<u>0.1MG</u>	N021137 005	Jun 06, 2003

PRESCRIPTION DRUG PRODUCT LIST

LEVOTHYROXINE SODIUM **

**See current Annual Edition, 1.8 Description of Special Situations, Levothyroxine Sodium

TABLET;ORAL

LEVOLET

	<u>AB3, AB4</u>	<u>0.1MG</u>			
-->	-->	<u>0.112MG</u>	N021137	006	Jun 06, 2003
	<u>AB1, AB2, AB3, AB4</u>				
-->	-->	<u>0.125MG</u>	N021137	007	Jun 06, 2003
	<u>AB1, AB2, AB3, AB4</u>				
-->	-->	<u>0.137MG</u>	N021137	008	Jun 06, 2003
	<u>AB1, AB2, AB3, AB4</u>				
-->	-->	<u>0.15MG</u>	N021137	009	Jun 06, 2003
	<u>AB1, AB2, AB3, AB4</u>				
-->	-->	<u>0.175MG</u>	N021137	010	Jun 06, 2003
	<u>AB1, AB2, AB3, AB4</u>				
-->	-->	<u>0.2MG</u>	N021137	011	Jun 06, 2003
	<u>AB1, AB2, AB3, AB4</u>				
-->	-->	<u>0.3MG</u>	N021137	012	Jun 06, 2003

LEVOTHYROXINE SODIUM

-->	ACCORD HLTHCARE	-->	<u>0.025MG</u>	A212399	001	Oct 19, 2020
			<u>AB1, AB2, AB3, AB4</u>			
-->		-->	<u>0.05MG</u>	A212399	002	Oct 19, 2020
			<u>AB1, AB2, AB3, AB4</u>			
-->		-->	<u>0.075MG</u>	A212399	003	Oct 19, 2020
			<u>AB1, AB2, AB3, AB4</u>			
-->		-->	<u>0.088MG</u>	A212399	004	Oct 19, 2020
			<u>AB1, AB2, AB3, AB4</u>			
-->		-->	<u>0.1MG</u>	A212399	005	Oct 19, 2020
			<u>AB1, AB2, AB3, AB4</u>			
-->		-->	<u>0.112MG</u>	A212399	006	Oct 19, 2020
			<u>AB1, AB2, AB3, AB4</u>			
-->		-->	<u>0.125MG</u>	A212399	007	Oct 19, 2020
			<u>AB1, AB2, AB3, AB4</u>			
-->		-->	<u>0.137MG</u>	A212399	008	Oct 19, 2020
			<u>AB1, AB2, AB3, AB4</u>			
-->		-->	<u>0.15MG</u>	A212399	009	Oct 19, 2020
			<u>AB1, AB2, AB3, AB4</u>			
-->		-->	<u>0.175MG</u>	A212399	010	Oct 19, 2020
			<u>AB1, AB2, AB3, AB4</u>			
-->		-->	<u>0.2MG</u>	A212399	011	Oct 19, 2020
			<u>AB1, AB2, AB3, AB4</u>			
-->		-->	<u>0.3MG</u>	A212399	012	Oct 19, 2020
			<u>AB1, AB2, AB3, AB4</u>			
-->	MYLAN	-->	<u>0.025MG</u>	A076187	001	Jun 05, 2002
			<u>AB1, AB2, AB3, AB4</u>			
-->		-->	<u>0.05MG</u>	A076187	002	Jun 05, 2002
			<u>AB1, AB2, AB3, AB4</u>			
-->		-->	<u>0.075MG</u>	A076187	003	Jun 05, 2002
			<u>AB1, AB2, AB3, AB4</u>			
-->		-->	<u>0.088MG</u>	A076187	004	Jun 05, 2002
			<u>AB1, AB2, AB3, AB4</u>			
-->		-->	<u>0.1MG</u>	A076187	005	Jun 05, 2002
			<u>AB1, AB2, AB3, AB4</u>			
-->		-->	<u>0.112MG</u>	A076187	006	Jun 05, 2002
			<u>AB1, AB2, AB3, AB4</u>			
-->		-->	<u>0.125MG</u>	A076187	007	Jun 05, 2002
			<u>AB1, AB2, AB3, AB4</u>			
-->		-->	<u>0.137MG</u>	A076187	012	Dec 13, 2006
			<u>AB1, AB2, AB3, AB4</u>			
-->		-->	<u>0.15MG</u>	A076187	008	Jun 05, 2002
			<u>AB1, AB2, AB3, AB4</u>			
-->		-->	<u>0.175MG</u>	A076187	009	Jun 05, 2002
			<u>AB1, AB2, AB3, AB4</u>			
-->		-->	<u>0.2MG</u>	A076187	010	Jun 05, 2002
			<u>AB1, AB2, AB3, AB4</u>			
-->	!	-->	<u>0.3MG</u>	A076187	011	Jun 05, 2002
			<u>AB1, AB2, AB3, AB4</u>			
-->	WATSON LABS TEVA	-->	<u>0.025MG</u>	A207588	001	May 10, 2022
			<u>AB1, AB2, AB3, AB4</u>			
-->		-->	<u>0.05MG</u>	A207588	002	May 10, 2022
			<u>AB1, AB2, AB3, AB4</u>			
-->		-->	<u>0.075MG</u>	A207588	003	May 10, 2022
			<u>AB1, AB2, AB3, AB4</u>			
-->		-->	<u>0.088MG</u>	A207588	004	May 10, 2022
			<u>AB1, AB2, AB3, AB4</u>			
-->		-->	<u>0.1MG</u>	A207588	005	May 10, 2022
			<u>AB1, AB2, AB3, AB4</u>			
-->		-->	<u>0.112MG</u>	A207588	006	May 10, 2022
			<u>AB1, AB2, AB3, AB4</u>			

PRESCRIPTION DRUG PRODUCT LIST

LEVOTHYROXINE SODIUM **

**See current Annual Edition, 1.8 Description of Special Situations, Levothyroxine Sodium

TABLET;ORAL

LEVOTHYROXINE SODIUM

-->		-->	<u>0.125MG</u>	A207588 007	May 10, 2022
		<u>AB1, AB2, AB3, AB4</u>			
-->		-->	<u>0.137MG</u>	A207588 008	May 10, 2022
		<u>AB1, AB2, AB3, AB4</u>			
-->		-->	<u>0.15MG</u>	A207588 009	May 10, 2022
		<u>AB1, AB2, AB3, AB4</u>			
-->		-->	<u>0.175MG</u>	A207588 010	May 10, 2022
		<u>AB1, AB2, AB3, AB4</u>			
-->		-->	<u>0.2MG</u>	A207588 011	May 10, 2022
		<u>AB1, AB2, AB3, AB4</u>			
-->		-->	<u>0.3MG</u>	A207588 012	May 10, 2022
		<u>AB1, AB2, AB3, AB4</u>			

THYRO-TABS

-->	+	ALVOGEN	-->	<u>0.025MG</u>	N021116 001	Oct 24, 2002
			<u>AB1, AB2, AB3, AB4</u>			
-->	+		-->	<u>0.05MG</u>	N021116 002	Oct 24, 2002
			<u>AB1, AB2, AB3, AB4</u>			
-->	+		-->	<u>0.075MG</u>	N021116 003	Oct 24, 2002
			<u>AB1, AB2, AB3, AB4</u>			
-->	+		-->	<u>0.088MG</u>	N021116 010	Oct 24, 2002
			<u>AB1, AB2, AB3, AB4</u>			
-->	+		-->	<u>0.1MG</u>	N021116 004	Oct 24, 2002
			<u>AB1, AB2, AB3, AB4</u>			
-->	+		-->	<u>0.112MG</u>	N021116 011	Oct 24, 2002
			<u>AB1, AB2, AB3, AB4</u>			
-->	+		-->	<u>0.125MG</u>	N021116 005	Oct 24, 2002
			<u>AB1, AB2, AB3, AB4</u>			
-->	+		-->	<u>0.137MG</u>	N021116 012	Dec 07, 2004
			<u>AB1, AB2, AB3, AB4</u>			
-->	+		-->	<u>0.15MG</u>	N021116 006	Oct 24, 2002
			<u>AB1, AB2, AB3, AB4</u>			
-->	+		-->	<u>0.175MG</u>	N021116 007	Oct 24, 2002
			<u>AB1, AB2, AB3, AB4</u>			
-->	+		-->	<u>0.2MG</u>	N021116 008	Oct 24, 2002
			<u>AB1, AB2, AB3, AB4</u>			
-->	+		-->	<u>0.3MG</u>	N021116 009	Oct 24, 2002
			<u>AB1, AB2, AB3, AB4</u>			

LEVOXYL

-->	+	KING PHARMS	-->	<u>AB1, AB3</u>	<u>0.025MG</u>	N021301 001	May 25, 2001
			-->	<u>AB1, AB3</u>	<u>0.05MG</u>	N021301 002	May 25, 2001
			-->	<u>AB1, AB3</u>	<u>0.075MG</u>	N021301 003	May 25, 2001
			-->	<u>AB1, AB3</u>	<u>0.088MG</u>	N021301 004	May 25, 2001
			-->	<u>AB1, AB3</u>	<u>0.1MG</u>	N021301 005	May 25, 2001
			-->	<u>AB1, AB3</u>	<u>0.112MG</u>	N021301 006	May 25, 2001
			-->	<u>AB1, AB3</u>	<u>0.125MG</u>	N021301 007	May 25, 2001
			-->	<u>AB1, AB3</u>	<u>0.137MG</u>	N021301 008	May 25, 2001
			-->	<u>AB1, AB3</u>	<u>0.15MG</u>	N021301 009	May 25, 2001
			-->	<u>AB1, AB3</u>	<u>0.175MG</u>	N021301 010	May 25, 2001
			-->	<u>AB1, AB3</u>	<u>0.2MG</u>	N021301 011	May 25, 2001

EUTHYROX

<u>AB2</u>	PROVELL	<u>0.025MG</u>	<u>N021292 001</u>	May 31, 2002
<u>AB2</u>		<u>0.05MG</u>	<u>N021292 002</u>	May 31, 2002
<u>AB2</u>		<u>0.075MG</u>	<u>N021292 003</u>	May 31, 2002
<u>AB2</u>		<u>0.088MG</u>	<u>N021292 004</u>	May 31, 2002
<u>AB2</u>		<u>0.1MG</u>	<u>N021292 005</u>	May 31, 2002
<u>AB2</u>		<u>0.112MG</u>	<u>N021292 006</u>	May 31, 2002
<u>AB2</u>		<u>0.125MG</u>	<u>N021292 007</u>	May 31, 2002
<u>AB2</u>		<u>0.137MG</u>	<u>N021292 008</u>	May 31, 2002
<u>AB2</u>		<u>0.15MG</u>	<u>N021292 009</u>	May 31, 2002
<u>AB2</u>		<u>0.175MG</u>	<u>N021292 010</u>	May 31, 2002
<u>AB2</u>		<u>0.2MG</u>	<u>N021292 011</u>	May 31, 2002

LEVOTHYROXINE SODIUM

<u>AB2</u>	AUROBINDO PHARMA	<u>0.137MG</u>	<u>A216414 001</u>	Jul 16, 2024
<u>AB2</u>		<u>0.15MG</u>	<u>A216414 002</u>	Jul 16, 2024
<u>AB2</u>		<u>0.175MG</u>	<u>A216414 003</u>	Jul 16, 2024
<u>AB2</u>		<u>0.2MG</u>	<u>A216414 004</u>	Jul 16, 2024
<u>AB2</u>		<u>0.3MG</u>	<u>A216414 005</u>	Jul 16, 2024
<u>AB4</u>	ASCENT PHARMS INC	<u>0.025MG</u>	<u>A215259 001</u>	Jan 18, 2023
<u>AB4</u>		<u>0.05MG</u>	<u>A215259 002</u>	Jan 18, 2023
<u>AB4</u>		<u>0.075MG</u>	<u>A215259 003</u>	Jan 18, 2023
<u>AB4</u>		<u>0.088MG</u>	<u>A215259 004</u>	Jan 18, 2023
<u>AB4</u>		<u>0.1MG</u>	<u>A215259 005</u>	Jan 18, 2023

PRESCRIPTION DRUG PRODUCT LIST

LEVOTHYROXINE SODIUM **

**See current Annual Edition, 1.8 Description of Special Situations, Levothyroxine Sodium

TABLET;ORAL

LEVOTHYROXINE SODIUM

<u>AB4</u>		<u>0.112MG</u>	<u>A215259 006</u>	Jan 18, 2023
<u>AB4</u>		<u>0.125MG</u>	<u>A215259 007</u>	Jan 18, 2023
<u>AB4</u>		<u>0.137MG</u>	<u>A215259 008</u>	Jan 18, 2023
<u>AB4</u>		<u>0.15MG</u>	<u>A215259 009</u>	Jan 18, 2023
<u>AB4</u>		<u>0.175MG</u>	<u>A215259 010</u>	Jan 18, 2023
<u>AB4</u>		<u>0.2MG</u>	<u>A215259 011</u>	Jan 18, 2023
<u>AB4</u>		<u>0.3MG</u>	<u>A215259 012</u>	Jan 18, 2023

LIDOCAINE

OINTMENT;TOPICAL

LIDOCAINE

<u>AT</u>	ALEMBIC	<u>5%</u>	<u>A211469 001</u>	Nov 23, 2018
<u>AT</u>	ALKEM LABS LTD	<u>5%</u>	<u>A207810 001</u>	Mar 10, 2017
<u>AT</u>	AMNEAL PHARMS	<u>5%</u>	<u>A206297 001</u>	Aug 07, 2015
<u>AT</u>	AUROBINDO PHARMA LTD	<u>5%</u>	<u>A217117 001</u>	Jun 18, 2024
<u>AT</u>	DR REDDYS	<u>5%</u>	<u>A208660 001</u>	Jan 05, 2021
<u>AT</u>	+! FOUGERA PHARMS INC	<u>5%</u>	<u>A080198 001</u>	
<u>AT</u>	GLENMARK PHARMS LTD	<u>5%</u>	<u>A206498 001</u>	Sep 09, 2016
<u>AT</u>	MACLEODS PHARMS LTD	<u>5%</u>	<u>A211697 001</u>	Mar 16, 2020
<u>AT</u>	QUAGEN	<u>5%</u>	<u>A212695 001</u>	Apr 20, 2021
<u>AT</u>	SEPTODONT INC	<u>5%</u>	<u>A040911 001</u>	May 23, 2011
<u>AT</u>	STRIDES PHARMA	<u>5%</u>	<u>A210958 001</u>	Dec 11, 2018
<u>AT</u>	TARO	<u>5%</u>	<u>A086724 001</u>	

PATCH;TOPICAL

LIDOCAINE

<u>AB</u>	ACTAVIS LABS UT INC	<u>5%</u>	<u>A200675 001</u>	Aug 23, 2012
<u>AB</u>	AMNEAL	<u>5%</u>	<u>A206463 001</u>	Aug 24, 2020
<u>AB</u>	IBSA	<u>5%</u>	<u>A209190 001</u>	Apr 30, 2020
<u>AB</u>	MYLAN TECHNOLOGIES	<u>5%</u>	<u>A202346 001</u>	Aug 07, 2015
<u>AB</u>	NAL PHARM	<u>5%</u>	<u>A205882 001</u>	Apr 29, 2021

LIDODERM

<u>AB</u>	+! TEIKOKU PHARMA USA	<u>5%</u>	<u>N020612 001</u>	Mar 19, 1999
	ZTLIDO			
	+! SCILEX PHARMS	<u>1.8%</u>	<u>N207962 001</u>	Feb 28, 2018

LIDOCAINE HYDROCHLORIDE

GEL;OPHTHALMIC

AKTEN

	+! THEA PHARMA	<u>3.5%</u>	<u>N022221 001</u>	Oct 07, 2008
--	----------------	-------------	--------------------	--------------

INJECTABLE;INJECTION

LIDOCAINE HYDROCHLORIDE

<u>AP</u>	AFAXYS	<u>1%</u>	<u>A215132 001</u>	Jun 09, 2022
<u>AP</u>	ASPIRO	<u>1%</u>	<u>A214336 001</u>	Nov 08, 2021
<u>AP</u>		<u>1%</u>	<u>A214339 001</u>	Nov 08, 2021
<u>AP</u>		<u>2%</u>	<u>A214336 002</u>	Dec 13, 2023
<u>AP</u>		<u>2%</u>	<u>A214339 002</u>	Nov 08, 2021
<u>AP</u>	B BRAUN MEDICAL INC	<u>1%</u>	<u>A208474 001</u>	Aug 03, 2018
<u>AP</u>	EUGIA PHARMA	<u>1%</u>	<u>A207182 001</u>	Oct 30, 2017
<u>AP</u>		<u>2%</u>	<u>A207182 002</u>	Oct 30, 2017
<u>AP</u>	FRESENIUS KABI USA	<u>1%</u>	<u>A088586 002</u>	
<u>AP</u>		<u>2%</u>	<u>A088586 003</u>	
<u>AP</u>	HOSPIRA	<u>0.5%</u>	<u>A088328 001</u>	May 17, 1984
<u>AP</u>	+!	<u>1%</u>	<u>A083158 001</u>	
<u>AP</u>		<u>1%</u>	<u>A088329 001</u>	May 17, 1984
<u>AP</u>		<u>2%</u>	<u>A040078 001</u>	Jun 23, 1995
<u>AP</u>	+!	<u>2%</u>	<u>A083158 002</u>	
<u>AP</u>		<u>2%</u>	<u>A088294 001</u>	May 17, 1984
<u>AP</u>	HUONS	<u>1%</u>	<u>A212821 001</u>	May 07, 2020
<u>AP</u>		<u>2%</u>	<u>A212821 002</u>	Jun 15, 2023
<u>AP</u>	+ INTL MEDICATION	<u>1%</u>	<u>A083173 001</u>	
<u>AP</u>		<u>2%</u>	<u>A083173 002</u>	
<u>AP</u>	MANKIND PHARMA	<u>1%</u>	<u>A217692 001</u>	Jun 16, 2023
<u>AP</u>		<u>1%</u>	<u>A217692 003</u>	Nov 29, 2023
<u>AP</u>		<u>1%</u>	<u>A217693 001</u>	Jul 11, 2023
<u>AP</u>		<u>2%</u>	<u>A217692 002</u>	Jun 16, 2023
<u>AP</u>		<u>2%</u>	<u>A217693 002</u>	Jul 11, 2023
<u>AP</u>	SINTETICA US	<u>1%</u>	<u>A214267 001</u>	Sep 19, 2022
<u>AP</u>		<u>2%</u>	<u>A214267 002</u>	Sep 19, 2022
<u>AP</u>		<u>4%</u>	<u>A214269 001</u>	May 05, 2022

PRESCRIPTION DRUG PRODUCT LIST

LIDOCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

LIDOCAINE HYDROCHLORIDE

<u>AP</u>	SPECTRA MDCL DEVICES	<u>1%</u>	<u>A208017 001</u>	Apr 18, 2018
<u>AP</u>	WEST-WARD PHARMS INT	<u>1%</u>	<u>A080407 001</u>	
<u>AP</u>		<u>2%</u>	<u>A080407 002</u>	

LIDOCAINE HYDROCHLORIDE 0.2% AND DEXTROSE 5% IN PLASTIC CONTAINER

<u>AP</u>	B BRAUN	<u>200MG/100ML</u>	<u>N019830 002</u>	Apr 08, 1992
<u>AP</u>	BAXTER HLTHCARE	<u>200MG/100ML</u>	<u>N018461 002</u>	

LIDOCAINE HYDROCHLORIDE 0.4% AND DEXTROSE 5% IN PLASTIC CONTAINER

<u>AP</u>	B BRAUN	<u>400MG/100ML</u>	<u>N019830 003</u>	Apr 08, 1992
<u>AP</u>	BAXTER HLTHCARE	<u>400MG/100ML</u>	<u>N018461 003</u>	

LIDOCAINE HYDROCHLORIDE 0.8% AND DEXTROSE 5% IN PLASTIC CONTAINER

<u>AP</u>	B BRAUN	<u>800MG/100ML</u>	<u>N019830 004</u>	Apr 08, 1992
<u>AP</u>	BAXTER HLTHCARE	<u>800MG/100ML</u>	<u>N018461 004</u>	Feb 22, 1982

LIDOCAINE HYDROCHLORIDE IN PLASTIC CONTAINER

<u>AP</u>	FRESENIUS KABI USA	<u>1%</u>	<u>A088586 001</u>	Jul 24, 1985
<u>AP</u>	HOSPIRA	<u>0.5%</u>	<u>A088325 001</u>	Jul 31, 1984
<u>AP</u>		<u>1%</u>	<u>A088299 001</u>	Jul 31, 1984
<u>AP</u>		<u>2%</u>	<u>A088327 001</u>	Jul 31, 1984

LIDOCAINE HYDROCHLORIDE PRESERVATIVE FREE

<u>AP</u>	EUGIA PHARMA	<u>1%</u>	<u>A203040 001</u>	Mar 14, 2013
<u>AP</u>		<u>1%</u>	<u>A203082 001</u>	Mar 14, 2013
<u>AP</u>		<u>2%</u>	<u>A203040 002</u>	Mar 14, 2013
<u>AP</u>		<u>2%</u>	<u>A203082 002</u>	Mar 14, 2013
<u>AP</u>	FRESENIUS KABI USA	<u>2%</u>	<u>N017584 001</u>	
<u>AP</u>		<u>4%</u>	<u>N017584 002</u>	
<u>AP</u>	HIKMA	<u>1%</u>	<u>A084625 001</u>	
<u>AP</u>		<u>2%</u>	<u>A084625 002</u>	
<u>AP</u>	HOSPIRA	<u>1%</u>	<u>A080408 001</u>	
<u>AP</u>		<u>1.5%</u>	<u>A080408 002</u>	
<u>AP</u>	!	<u>4%</u>	<u>A088295 001</u>	May 17, 1984

LIDOCAINE HYDROCHLORIDE PRESERVATIVE FREE IN PLASTIC CONTAINER

<u>AP</u>	HOSPIRA	<u>1%</u>	<u>A040302 001</u>	Sep 28, 1998
<u>AP</u>		<u>2%</u>	<u>A040302 002</u>	Sep 28, 1998

XYLOCAINE

<u>AP</u>	+!	FRESENIUS KABI USA	<u>0.5%</u>	<u>N006488 008</u>
<u>AP</u>	+!		<u>1%</u>	<u>N006488 007</u>
<u>AP</u>	+!		<u>1.5%</u>	<u>N006488 010</u>
<u>AP</u>	+!		<u>2%</u>	<u>N006488 002</u>

INJECTABLE; SPINAL

LIDOCAINE HYDROCHLORIDE 5% AND DEXTROSE 7.5%

+! HOSPIRA 5% A083914 001

JELLY; TOPICAL

GLYDO

<u>AT</u>	!	SAGENT PHARMS INC	<u>2%</u>	<u>A201094 001</u>	Apr 28, 2014
-----------	---	-------------------	-----------	--------------------	--------------

LIDOCAINE HYDROCHLORIDE

<u>AT</u>		INTL MEDICATION	<u>2%</u>	<u>A086283 001</u>	
<u>AT</u>		SENTISS	<u>2%</u>	<u>A040433 001</u>	Feb 12, 2003

SOLUTION; ORAL

LIDOCAINE HYDROCHLORIDE

<u>AT</u>	!	RUBICON	<u>2%</u>	<u>A216780 001</u>	Mar 28, 2023
<u>AT</u>		WOCKHARDT BIO AG	<u>2%</u>	<u>A087872 001</u>	Nov 18, 1982

LIDOCAINE HYDROCHLORIDE VISCOUS

<u>AT</u>		CHARTWELL MOLECULAR	<u>2%</u>	<u>A040708 001</u>	Feb 27, 2007
<u>AT</u>		PAI HOLDINGS PHARM	<u>2%</u>	<u>A218138 001</u>	Feb 20, 2024

LIDOCAINE VISCOUS

<u>AT</u>		HIKMA	<u>2%</u>	<u>A088802 001</u>	Apr 26, 1985
-----------	--	-------	-----------	--------------------	--------------

SOLUTION; TOPICAL

LARYNG-O-JET KIT

<u>AT</u>		INTL MEDICATION	<u>4%</u>	<u>A086364 001</u>	
-----------	--	-----------------	-----------	--------------------	--

LIDOCAINE HYDROCHLORIDE

<u>AT</u>	!	HIKMA	<u>4%</u>	<u>A088803 001</u>	Apr 03, 1985
<u>AT</u>		LANNETT CO INC	<u>4%</u>	<u>A040710 001</u>	Feb 27, 2007
<u>AT</u>		NOVITIUM PHARMA	<u>4%</u>	<u>A216250 001</u>	Mar 23, 2022
<u>AT</u>		PAI HOLDINGS PHARM	<u>4%</u>	<u>A204494 001</u>	Mar 12, 2014
<u>AT</u>		TARO	<u>4%</u>	<u>A218182 001</u>	Dec 07, 2023
<u>AT</u>		THE J MOLNER	<u>4%</u>	<u>A218411 001</u>	Apr 29, 2024
<u>AT</u>		WOCKHARDT BIO AG	<u>4%</u>	<u>A087881 001</u>	Nov 18, 1982

PRESCRIPTION DRUG PRODUCT LIST

LIDOCAINE HYDROCHLORIDE

SYSTEM; INTRADERMAL

ZINGO

POWDER PHARMS

0.5MG

N022114 001 Aug 16, 2007

LIDOCAINE; PRILUCAINE

CREAM; TOPICAL

LIDOCAINE AND PRILUCAINE**AB** ALEMBIC2.5%;2.5%A213923 001 Apr 08, 2022**AB** ENCUBE2.5%;2.5%A076320 001 Aug 27, 2003**AB** ! FOUGERA PHARMS2.5%;2.5%A076453 001 Aug 18, 2003**AB** PADAGIS US2.5%;2.5%A212482 001 Jul 27, 2021**AB** PAI HOLDINGS PHARM2.5%;2.5%A205887 001 Jun 29, 2018**AB** ZYDUS LIFESCIENCES2.5%;2.5%A219120 001 Dec 17, 2024

GEL; PERIODONTAL

ORAQIX

+! DENTSPLY PHARM

2.5%;2.5%

N021451 001 Dec 19, 2003

LIFITEGRAST

SOLUTION/DROPS; OPHTHALMIC

XIIDRA

+! BAUSCH AND LOMB INC

5%

N208073 001 Jul 11, 2016

LINACLOTIDE

CAPSULE; ORAL

LINZESS

+! ABBVIE

72MCG

N202811 003 Jan 25, 2017

+!

145MCG

N202811 001 Aug 30, 2012

+

290MCG

N202811 002 Aug 30, 2012

LINAGLIPTIN

TABLET; ORAL

LINAGLIPTIN**AB** INVAGEN PHARMS5MGA208423 001 Sep 03, 2024**AB** SUNSHINE5MGA208335 001 Aug 31, 2021TRADJENTA**AB** +! BOEHRINGER5MGN201280 001 May 02, 2011

INGELHEIM

LINAGLIPTIN; METFORMIN HYDROCHLORIDE

TABLET; ORAL

JENTADUETO**AB** + BOEHRINGER2.5MG;500MGN201281 001 Jan 30, 2012

INGELHEIM

AB +2.5MG;850MGN201281 002 Jan 30, 2012**AB** +!2.5MG;1GMN201281 003 Jan 30, 2012LINAGLIPTIN AND METFORMIN HYDROCHLORIDE**AB** SUNSHINE2.5MG;500MGA208336 001 Aug 30, 2021**AB**2.5MG;850MGA208336 002 Aug 30, 2021**AB**2.5MG;1GMA208336 003 Aug 30, 2021

TABLET, EXTENDED RELEASE; ORAL

JENTADUETO XR

+ BOEHRINGER

2.5MG;1GM

N208026 001 May 27, 2016

INGELHEIM

+!

5MG;1GM

N208026 002 May 27, 2016

LINCOMYCIN HYDROCHLORIDE

INJECTABLE; INJECTION

LINCOCIN**AP** +! PFIZEREQ 300MG BASE/MLN050317 001LINCOMYCIN HYDROCHLORIDE**AP** ARTHUR GRPEQ 300MG BASE/MLA212770 001 Mar 12, 2021**AP** GLAND PHARMA LTDEQ 300MG BASE/MLA215657 001 Sep 26, 2022**AP** MICRO LABSEQ 300MG BASE/MLA215082 001 Nov 08, 2021**AP** XGEN PHARMSEQ 300MG BASE/MLA201746 001 Jun 04, 2015LINEZOLID

FOR SUSPENSION; ORAL

LINEZOLID**AB** HETERO LABS100MG/5MLA211813 001 Oct 31, 2022**AB** HIKMA100MG/5MLA200068 001 Jun 03, 2015ZYVOX**AB** +! PFIZER100MG/5MLN021132 001 Apr 18, 2000

SOLUTION; INTRAVENOUS

LINEZOLID**AP** EUGIA PHARMA600MG/300ML (2MG/ML)A206917 001 Aug 04, 2016

PRESCRIPTION DRUG PRODUCT LIST

LINEZOLID

SOLUTION; INTRAVENOUS

LINEZOLID

<u>AP</u>	FRENIUS KABI USA	<u>600MG/300ML (2MG/ML)</u>	<u>A204764 001</u>	Mar 15, 2016
<u>AP</u>	HIKMA	<u>600MG/300ML (2MG/ML)</u>	<u>A206454 001</u>	Aug 22, 2022
<u>AP</u>	HQ SPCLT PHARMA	<u>200MG/100ML (2MG/ML)</u>	<u>A207001 001</u>	Jul 07, 2017
<u>AP</u>		<u>600MG/300ML (2MG/ML)</u>	<u>A207001 002</u>	Jul 07, 2017
<u>AP</u>	MYLAN LABS LTD	<u>200MG/100ML (2MG/ML)</u>	<u>A205154 001</u>	Dec 06, 2017
<u>AP</u>		<u>600MG/300ML (2MG/ML)</u>	<u>A205154 002</u>	Dec 06, 2017
<u>AP</u>	NANG KUANG PHARM CO	<u>200MG/100ML (2MG/ML)</u>	<u>A207354 001</u>	Dec 20, 2016
<u>AP</u>		<u>600MG/300ML (2MG/ML)</u>	<u>A207354 002</u>	Dec 20, 2016
<u>AP</u>	SAGENT PHARMS INC	<u>200MG/100ML (2MG/ML)</u>	<u>A204696 001</u>	Mar 02, 2017
<u>AP</u>		<u>600MG/300ML (2MG/ML)</u>	<u>A204696 002</u>	Mar 02, 2017
<u>AP</u>	SANDOZ	<u>200MG/100ML (2MG/ML)</u>	<u>A200904 001</u>	Jul 16, 2015
<u>AP</u>		<u>600MG/300ML (2MG/ML)</u>	<u>A200904 002</u>	Jul 16, 2015

ZYVOX

<u>AP</u>	+	PFIZER	<u>200MG/100ML (2MG/ML)</u>	<u>N021131 001</u>	Apr 18, 2000	
<u>AP</u>	+	!	<u>600MG/300ML (2MG/ML)</u>	<u>N021131 003</u>	Apr 18, 2000	
		LINEZOLID IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER				
		+	HOSPIRA	<u>600MG/300ML (2MG/ML)</u>	<u>N206473 001</u>	Jun 18, 2015

TABLET; ORAL

LINEZOLID

<u>AB</u>	ALEMBIC	<u>600MG</u>	<u>A205233 001</u>	Dec 21, 2015
<u>AB</u>	ALKEM LABS LTD	<u>600MG</u>	<u>A205517 001</u>	Dec 21, 2015
<u>AB</u>	CHARTWELL RX	<u>600MG</u>	<u>A210702 001</u>	Apr 25, 2019
<u>AB</u>	GLENMARK SPECLT	<u>600MG</u>	<u>A078987 001</u>	Dec 21, 2015
<u>AB</u>	HETERO LABS LTD V	<u>600MG</u>	<u>A204239 001</u>	Dec 21, 2015
<u>AB</u>	MACLEODS PHARMS LTD	<u>600MG</u>	<u>A210378 001</u>	Oct 10, 2024
<u>AB</u>	NOVEL LABS INC	<u>600MG</u>	<u>A207526 001</u>	Aug 22, 2016
<u>AB</u>	RISING	<u>600MG</u>	<u>A078845 001</u>	Dec 21, 2015
<u>AB</u>	ZYDUS PHARMS	<u>600MG</u>	<u>A206097 001</u>	Feb 22, 2017

ZYVOX

<u>AB</u>	+	!	PFIZER	<u>600MG</u>	<u>N021130 002</u>	Apr 18, 2000
-----------	---	---	--------	--------------	--------------------	--------------

LIOTHYRONINE SODIUM

INJECTABLE; INJECTION

LIOTHYRONINE SODIUM

!

XGEN PHARMS

EQ 0.01MG BASE/ML

A076923 001

Aug 17, 2005

TABLET; ORAL

CYTOMEL

<u>AB</u>	+	KING PHARMS	<u>EQ 0.005MG BASE</u>	<u>N010379 001</u>
<u>AB</u>	+		<u>EQ 0.025MG BASE</u>	<u>N010379 002</u>
<u>AB</u>	+		<u>EQ 0.05MG BASE</u>	<u>N010379 003</u>

LIOTHYRONINE SODIUM

<u>AB</u>	BIOCON PHARMA	<u>EQ 0.005MG BASE</u>	<u>A218070 001</u>	Feb 06, 2024
<u>AB</u>		<u>EQ 0.025MG BASE</u>	<u>A218070 002</u>	Feb 06, 2024
<u>AB</u>		<u>EQ 0.05MG BASE</u>	<u>A218070 003</u>	Feb 06, 2024
<u>AB</u>	DR REDDYS LABS SA	<u>EQ 0.005MG BASE</u>	<u>A090097 001</u>	Mar 20, 2009
<u>AB</u>		<u>EQ 0.025MG BASE</u>	<u>A090097 002</u>	Mar 20, 2009
<u>AB</u>		<u>EQ 0.05MG BASE</u>	<u>A090097 003</u>	Mar 20, 2009
<u>AB</u>	SIGMAPHARM LABS LLC	<u>EQ 0.005MG BASE</u>	<u>A200295 001</u>	Nov 29, 2012
<u>AB</u>		<u>EQ 0.025MG BASE</u>	<u>A200295 002</u>	Nov 29, 2012
<u>AB</u>	!	<u>EQ 0.05MG BASE</u>	<u>A200295 003</u>	Nov 29, 2012
<u>AB</u>	SUN PHARM	<u>EQ 0.005MG BASE</u>	<u>A091382 001</u>	Apr 20, 2016
<u>AB</u>		<u>EQ 0.025MG BASE</u>	<u>A091382 002</u>	Apr 20, 2016
<u>AB</u>		<u>EQ 0.05MG BASE</u>	<u>A091382 003</u>	Apr 20, 2016
<u>AB</u>	TEVA PHARMS USA	<u>EQ 0.005MG BASE</u>	<u>A211510 001</u>	Oct 26, 2018
<u>AB</u>		<u>EQ 0.025MG BASE</u>	<u>A211510 002</u>	Oct 26, 2018
<u>AB</u>		<u>EQ 0.05MG BASE</u>	<u>A211510 003</u>	Oct 26, 2018
<u>AB</u>	ZYDUS LIFESCIENCES	<u>EQ 0.005MG BASE</u>	<u>A214803 001</u>	Jan 22, 2021
<u>AB</u>		<u>EQ 0.025MG BASE</u>	<u>A214803 002</u>	Jan 22, 2021
<u>AB</u>		<u>EQ 0.05MG BASE</u>	<u>A214803 003</u>	Jan 22, 2021

LIRAGLUTIDE

SOLUTION; SUBCUTANEOUS

LIRAGLUTIDE

<u>AP</u>	HIKMA	<u>18MG/3ML (6MG/ML)</u>	<u>A215503 001</u>	Dec 23, 2024		
<u>AP</u>	+	!	NOVO NORDISK INC	<u>18MG/3ML (6MG/ML)</u>	<u>N022341 001</u>	Jan 25, 2010
	SAXENDA					
	+	!	NOVO	<u>18MG/3ML (6MG/ML)</u>	<u>N206321 001</u>	Dec 23, 2014

PRESCRIPTION DRUG PRODUCT LIST

LISDEXAMFETAMINE DIMESYLATE

CAPSULE; ORAL

LISDEXAMFETAMINE DIMESYLATE

<u>AB</u>	ACTAVIS ELIZABETH	<u>10MG</u>	<u>A202802 001</u>	Aug 25, 2023
<u>AB</u>		<u>20MG</u>	<u>A202802 002</u>	Aug 25, 2023
<u>AB</u>		<u>30MG</u>	<u>A202802 003</u>	Aug 25, 2023
<u>AB</u>		<u>40MG</u>	<u>A202802 004</u>	Aug 25, 2023
<u>AB</u>		<u>50MG</u>	<u>A202802 005</u>	Aug 25, 2023
<u>AB</u>		<u>60MG</u>	<u>A202802 006</u>	Aug 25, 2023
<u>AB</u>		<u>70MG</u>	<u>A202802 007</u>	Aug 25, 2023
<u>AB</u>	ALKEM LABS LTD	<u>10MG</u>	<u>A217194 001</u>	Aug 25, 2023
<u>AB</u>		<u>20MG</u>	<u>A217194 002</u>	Aug 25, 2023
<u>AB</u>		<u>30MG</u>	<u>A217194 003</u>	Aug 25, 2023
<u>AB</u>		<u>40MG</u>	<u>A217194 004</u>	Aug 25, 2023
<u>AB</u>		<u>50MG</u>	<u>A217194 005</u>	Aug 25, 2023
<u>AB</u>		<u>60MG</u>	<u>A217194 006</u>	Aug 25, 2023
<u>AB</u>		<u>70MG</u>	<u>A217194 007</u>	Aug 25, 2023
<u>AB</u>	AMNEAL	<u>20MG</u>	<u>A202830 001</u>	Aug 25, 2023
<u>AB</u>		<u>30MG</u>	<u>A202830 002</u>	Aug 25, 2023
<u>AB</u>		<u>40MG</u>	<u>A202830 003</u>	Aug 25, 2023
<u>AB</u>		<u>50MG</u>	<u>A202830 004</u>	Aug 25, 2023
<u>AB</u>		<u>60MG</u>	<u>A202830 005</u>	Aug 25, 2023
<u>AB</u>		<u>70MG</u>	<u>A202830 006</u>	Aug 25, 2023
<u>AB</u>	APOTEX	<u>10MG</u>	<u>A216944 001</u>	Aug 25, 2023
<u>AB</u>		<u>20MG</u>	<u>A216944 002</u>	Aug 25, 2023
<u>AB</u>		<u>30MG</u>	<u>A216944 003</u>	Aug 25, 2023
<u>AB</u>		<u>40MG</u>	<u>A216944 004</u>	Aug 25, 2023
<u>AB</u>		<u>50MG</u>	<u>A216944 005</u>	Aug 25, 2023
<u>AB</u>		<u>60MG</u>	<u>A216944 006</u>	Aug 25, 2023
<u>AB</u>		<u>70MG</u>	<u>A216944 007</u>	Aug 25, 2023
<u>AB</u>	ASCENT PHARMS INC	<u>10MG</u>	<u>A217442 001</u>	Aug 25, 2023
<u>AB</u>		<u>20MG</u>	<u>A217442 002</u>	Aug 25, 2023
<u>AB</u>		<u>30MG</u>	<u>A217442 003</u>	Aug 25, 2023
<u>AB</u>		<u>40MG</u>	<u>A217442 004</u>	Aug 25, 2023
<u>AB</u>		<u>50MG</u>	<u>A217442 005</u>	Aug 25, 2023
<u>AB</u>		<u>60MG</u>	<u>A217442 006</u>	Aug 25, 2023
<u>AB</u>		<u>70MG</u>	<u>A217442 007</u>	Aug 25, 2023
<u>AB</u>	ELITE LABS INC	<u>10MG</u>	<u>A218604 001</u>	Nov 15, 2024
<u>AB</u>		<u>20MG</u>	<u>A218604 002</u>	Nov 15, 2024
<u>AB</u>		<u>30MG</u>	<u>A218604 003</u>	Nov 15, 2024
<u>AB</u>		<u>40MG</u>	<u>A218604 004</u>	Nov 15, 2024
<u>AB</u>		<u>50MG</u>	<u>A218604 005</u>	Nov 15, 2024
<u>AB</u>		<u>60MG</u>	<u>A218604 006</u>	Nov 15, 2024
<u>AB</u>		<u>70MG</u>	<u>A218604 007</u>	Nov 15, 2024
<u>AB</u>	HIKMA	<u>10MG</u>	<u>A202827 007</u>	Jan 17, 2024
<u>AB</u>		<u>20MG</u>	<u>A202827 001</u>	Aug 25, 2023
<u>AB</u>		<u>30MG</u>	<u>A202827 002</u>	Aug 25, 2023
<u>AB</u>		<u>40MG</u>	<u>A202827 003</u>	Aug 25, 2023
<u>AB</u>		<u>50MG</u>	<u>A202827 004</u>	Aug 25, 2023
<u>AB</u>		<u>60MG</u>	<u>A202827 005</u>	Aug 25, 2023
<u>AB</u>		<u>70MG</u>	<u>A202827 006</u>	Aug 25, 2023
<u>AB</u>	LANNETT CO INC	<u>10MG</u>	<u>A215802 001</u>	Aug 25, 2023
<u>AB</u>		<u>20MG</u>	<u>A215802 002</u>	Aug 25, 2023
<u>AB</u>		<u>30MG</u>	<u>A215802 003</u>	Aug 25, 2023
<u>AB</u>		<u>40MG</u>	<u>A215802 004</u>	Aug 25, 2023
<u>AB</u>		<u>50MG</u>	<u>A215802 005</u>	Aug 25, 2023
<u>AB</u>		<u>60MG</u>	<u>A215802 006</u>	Aug 25, 2023
<u>AB</u>		<u>70MG</u>	<u>A215802 007</u>	Aug 25, 2023
<u>AB</u>	MYLAN	<u>10MG</u>	<u>A202835 001</u>	Aug 25, 2023
<u>AB</u>		<u>20MG</u>	<u>A202835 002</u>	Aug 25, 2023
<u>AB</u>		<u>30MG</u>	<u>A202835 003</u>	Aug 25, 2023
<u>AB</u>		<u>40MG</u>	<u>A202835 004</u>	Aug 25, 2023
<u>AB</u>		<u>50MG</u>	<u>A202835 005</u>	Aug 25, 2023
<u>AB</u>		<u>60MG</u>	<u>A202835 006</u>	Aug 25, 2023
<u>AB</u>		<u>70MG</u>	<u>A202835 007</u>	Aug 25, 2023
<u>AB</u>	NORWICH	<u>10MG</u>	<u>A214547 001</u>	Aug 25, 2023
<u>AB</u>		<u>20MG</u>	<u>A214547 002</u>	Aug 25, 2023
<u>AB</u>		<u>30MG</u>	<u>A214547 003</u>	Aug 25, 2023
<u>AB</u>		<u>40MG</u>	<u>A214547 004</u>	Aug 25, 2023
<u>AB</u>		<u>50MG</u>	<u>A214547 005</u>	Aug 25, 2023
<u>AB</u>		<u>60MG</u>	<u>A214547 006</u>	Aug 25, 2023
<u>AB</u>		<u>70MG</u>	<u>A214547 007</u>	Aug 25, 2023

PRESCRIPTION DRUG PRODUCT LIST

LISDEXAMFETAMINE DIMESYLATE

CAPSULE;ORAL

LISDEXAMFETAMINE DIMESYLATE

<u>AB</u>	PRINSTON INC	<u>10MG</u>	<u>A216266 001</u>	Aug 25, 2023
<u>AB</u>		<u>20MG</u>	<u>A216266 002</u>	Aug 25, 2023
<u>AB</u>		<u>30MG</u>	<u>A216266 003</u>	Aug 25, 2023
<u>AB</u>		<u>40MG</u>	<u>A216266 004</u>	Aug 25, 2023
<u>AB</u>		<u>50MG</u>	<u>A216266 005</u>	Aug 25, 2023
<u>AB</u>		<u>60MG</u>	<u>A216266 006</u>	Aug 25, 2023
<u>AB</u>		<u>70MG</u>	<u>A216266 007</u>	Aug 25, 2023
<u>AB</u>	RHODES PHARMS	<u>10MG</u>	<u>A215330 001</u>	Aug 25, 2023
<u>AB</u>		<u>20MG</u>	<u>A215330 002</u>	Aug 25, 2023
<u>AB</u>		<u>30MG</u>	<u>A215330 003</u>	Aug 25, 2023
<u>AB</u>		<u>40MG</u>	<u>A215330 004</u>	Aug 25, 2023
<u>AB</u>		<u>50MG</u>	<u>A215330 005</u>	Aug 25, 2023
<u>AB</u>		<u>60MG</u>	<u>A215330 006</u>	Aug 25, 2023
<u>AB</u>		<u>70MG</u>	<u>A215330 007</u>	Aug 25, 2023
<u>AB</u>	SANDOZ	<u>10MG</u>	<u>A202836 007</u>	Dec 26, 2024
<u>AB</u>	SPECGX LLC	<u>10MG</u>	<u>A211840 001</u>	Aug 25, 2023
<u>AB</u>		<u>20MG</u>	<u>A211840 002</u>	Aug 25, 2023
<u>AB</u>		<u>30MG</u>	<u>A211840 003</u>	Aug 25, 2023
<u>AB</u>		<u>40MG</u>	<u>A211840 004</u>	Aug 25, 2023
<u>AB</u>		<u>50MG</u>	<u>A211840 005</u>	Aug 25, 2023
<u>AB</u>		<u>60MG</u>	<u>A211840 006</u>	Aug 25, 2023
<u>AB</u>		<u>70MG</u>	<u>A211840 007</u>	Aug 25, 2023
<u>AB</u>	SUN PHARM INDS INC	<u>10MG</u>	<u>A214484 001</u>	Aug 25, 2023
<u>AB</u>		<u>20MG</u>	<u>A214484 002</u>	Aug 25, 2023
<u>AB</u>		<u>30MG</u>	<u>A214484 003</u>	Aug 25, 2023
<u>AB</u>		<u>40MG</u>	<u>A214484 004</u>	Aug 25, 2023
<u>AB</u>		<u>50MG</u>	<u>A214484 005</u>	Aug 25, 2023
<u>AB</u>		<u>60MG</u>	<u>A214484 006</u>	Aug 25, 2023
<u>AB</u>		<u>70MG</u>	<u>A214484 007</u>	Aug 25, 2023
<u>VYVANSE</u>				
<u>AB</u>	+ TAKEDA PHARMS USA	<u>10MG</u>	<u>N021977 007</u>	Oct 30, 2014
<u>AB</u>	+	<u>20MG</u>	<u>N021977 004</u>	Dec 10, 2007
<u>AB</u>	+	<u>30MG</u>	<u>N021977 001</u>	Feb 23, 2007
<u>AB</u>	+	<u>40MG</u>	<u>N021977 005</u>	Dec 10, 2007
<u>AB</u>	+	<u>50MG</u>	<u>N021977 002</u>	Feb 23, 2007
<u>AB</u>	+	<u>60MG</u>	<u>N021977 006</u>	Dec 10, 2007
<u>AB</u>	+	<u>70MG</u>	<u>N021977 003</u>	Feb 23, 2007

TABLET, CHEWABLE;ORAL

LISDEXAMFETAMINE DIMESYLATE

<u>AB</u>	ASCENT PHARMS INC	<u>10MG</u>	<u>A217068 001</u>	Aug 25, 2023
<u>AB</u>		<u>20MG</u>	<u>A217068 002</u>	Aug 25, 2023
<u>AB</u>		<u>30MG</u>	<u>A217068 003</u>	Aug 25, 2023
<u>AB</u>		<u>40MG</u>	<u>A217068 004</u>	Aug 25, 2023
<u>AB</u>		<u>50MG</u>	<u>A217068 005</u>	Aug 25, 2023
<u>AB</u>		<u>60MG</u>	<u>A217068 006</u>	Aug 25, 2023
<u>AB</u>	GRANULES	<u>10MG</u>	<u>A219258 001</u>	Dec 16, 2024
<u>AB</u>		<u>20MG</u>	<u>A219258 002</u>	Dec 16, 2024
<u>AB</u>		<u>30MG</u>	<u>A219258 003</u>	Dec 16, 2024
<u>AB</u>		<u>40MG</u>	<u>A219258 004</u>	Dec 16, 2024
<u>AB</u>		<u>50MG</u>	<u>A219258 005</u>	Dec 16, 2024
<u>AB</u>		<u>60MG</u>	<u>A219258 006</u>	Dec 16, 2024
<u>AB</u>	MSN	<u>10MG</u>	<u>A218306 001</u>	Feb 02, 2024
<u>AB</u>		<u>20MG</u>	<u>A218306 002</u>	Feb 02, 2024
<u>AB</u>		<u>30MG</u>	<u>A218306 003</u>	Feb 02, 2024
<u>AB</u>		<u>40MG</u>	<u>A218306 004</u>	Feb 02, 2024
<u>AB</u>		<u>50MG</u>	<u>A218306 005</u>	Feb 02, 2024
<u>AB</u>		<u>60MG</u>	<u>A218306 006</u>	Feb 02, 2024
<u>AB</u>	SPECGX LLC	<u>10MG</u>	<u>A218850 001</u>	Dec 17, 2024
<u>AB</u>		<u>20MG</u>	<u>A218850 002</u>	Dec 17, 2024
<u>AB</u>		<u>30MG</u>	<u>A218850 003</u>	Dec 17, 2024
<u>AB</u>		<u>40MG</u>	<u>A218850 004</u>	Dec 17, 2024
<u>AB</u>		<u>50MG</u>	<u>A218850 005</u>	Dec 17, 2024
<u>AB</u>		<u>60MG</u>	<u>A218850 006</u>	Dec 17, 2024
<u>AB</u>	SUN PHARM INDS INC	<u>10MG</u>	<u>A214134 001</u>	Aug 25, 2023
<u>AB</u>		<u>20MG</u>	<u>A214134 002</u>	Aug 25, 2023
<u>AB</u>		<u>30MG</u>	<u>A214134 003</u>	Aug 25, 2023
<u>AB</u>		<u>40MG</u>	<u>A214134 004</u>	Aug 25, 2023
<u>AB</u>		<u>50MG</u>	<u>A214134 005</u>	Aug 25, 2023
<u>AB</u>		<u>60MG</u>	<u>A214134 006</u>	Aug 25, 2023

PRESCRIPTION DRUG PRODUCT LIST

LISDEXAMFETAMINE DIMESYLATE

TABLET, CHEWABLE;ORAL

LISDEXAMFETAMINE DIMESYLATE

<u>AB</u>	TEVA PHARMS	<u>10MG</u>	<u>A215415 001</u>	Aug 25, 2023
<u>AB</u>		<u>20MG</u>	<u>A215415 002</u>	Aug 25, 2023
<u>AB</u>		<u>30MG</u>	<u>A215415 003</u>	Aug 25, 2023
<u>AB</u>		<u>40MG</u>	<u>A215415 004</u>	Aug 25, 2023
<u>AB</u>		<u>50MG</u>	<u>A215415 005</u>	Aug 25, 2023
<u>AB</u>		<u>60MG</u>	<u>A215415 006</u>	Aug 25, 2023

VYVANSE

<u>AB</u>	+	TAKEDA PHARMS USA	<u>10MG</u>	<u>N208510 001</u>	Jan 28, 2017
<u>AB</u>	+		<u>20MG</u>	<u>N208510 002</u>	Jan 28, 2017
<u>AB</u>	+		<u>30MG</u>	<u>N208510 003</u>	Jan 28, 2017
<u>AB</u>	+		<u>40MG</u>	<u>N208510 004</u>	Jan 28, 2017
<u>AB</u>	+		<u>50MG</u>	<u>N208510 005</u>	Jan 28, 2017
<u>AB</u>	+	!	<u>60MG</u>	<u>N208510 006</u>	Jan 28, 2017

LISINOPRIL

SOLUTION;ORAL

QBRELIS

+! AZURITY

1MG/ML

N208401 001 Jul 29, 2016

TABLET;ORAL

LISINOPRIL

<u>AB</u>		ASCENT PHARMS INC	<u>2.5MG</u>	<u>A075903 001</u>	Jul 01, 2002
<u>AB</u>			<u>5MG</u>	<u>A075903 002</u>	Jul 01, 2002
<u>AB</u>			<u>10MG</u>	<u>A075903 003</u>	Jul 01, 2002
<u>AB</u>			<u>20MG</u>	<u>A075903 004</u>	Jul 01, 2002
<u>AB</u>			<u>30MG</u>	<u>A075903 005</u>	Jul 01, 2002
<u>AB</u>			<u>40MG</u>	<u>A075903 006</u>	Jul 01, 2002
<u>AB</u>		AUROBINDO	<u>2.5MG</u>	<u>A077622 001</u>	Feb 22, 2006
<u>AB</u>			<u>5MG</u>	<u>A077622 002</u>	Feb 22, 2006
<u>AB</u>			<u>10MG</u>	<u>A077622 003</u>	Feb 22, 2006
<u>AB</u>			<u>20MG</u>	<u>A077622 004</u>	Feb 22, 2006
<u>AB</u>			<u>30MG</u>	<u>A077622 005</u>	Feb 22, 2006
<u>AB</u>			<u>40MG</u>	<u>A077622 006</u>	Feb 22, 2006
<u>AB</u>		CHARTWELL RX	<u>2.5MG</u>	<u>A075994 001</u>	Jul 01, 2002
<u>AB</u>			<u>5MG</u>	<u>A075994 002</u>	Jul 01, 2002
<u>AB</u>			<u>10MG</u>	<u>A075994 003</u>	Jul 01, 2002
<u>AB</u>			<u>20MG</u>	<u>A075994 004</u>	Jul 01, 2002
<u>AB</u>			<u>30MG</u>	<u>A075994 005</u>	Jul 01, 2002
<u>AB</u>			<u>40MG</u>	<u>A075994 006</u>	Jul 01, 2002
<u>AB</u>		COREPHARMA	<u>2.5MG</u>	<u>A076102 001</u>	Sep 30, 2002
<u>AB</u>			<u>5MG</u>	<u>A076102 002</u>	Sep 30, 2002
<u>AB</u>			<u>10MG</u>	<u>A076102 003</u>	Sep 30, 2002
<u>AB</u>			<u>20MG</u>	<u>A076102 004</u>	Sep 30, 2002
<u>AB</u>			<u>30MG</u>	<u>A076102 005</u>	Sep 30, 2002
<u>AB</u>			<u>40MG</u>	<u>A076102 006</u>	Sep 30, 2002
<u>AB</u>		INVAGEN PHARMS	<u>2.5MG</u>	<u>A203508 001</u>	Oct 29, 2013
<u>AB</u>			<u>5MG</u>	<u>A203508 002</u>	Oct 29, 2013
<u>AB</u>			<u>10MG</u>	<u>A203508 003</u>	Oct 29, 2013
<u>AB</u>			<u>20MG</u>	<u>A203508 004</u>	Oct 29, 2013
<u>AB</u>			<u>30MG</u>	<u>A203508 005</u>	Oct 29, 2013
<u>AB</u>			<u>40MG</u>	<u>A203508 006</u>	Oct 29, 2013
<u>AB</u>		LUPIN	<u>2.5MG</u>	<u>A077321 001</u>	Sep 09, 2005
<u>AB</u>			<u>5MG</u>	<u>A077321 002</u>	Sep 09, 2005
<u>AB</u>			<u>10MG</u>	<u>A077321 003</u>	Sep 09, 2005
<u>AB</u>			<u>20MG</u>	<u>A077321 004</u>	Sep 09, 2005
<u>AB</u>			<u>30MG</u>	<u>A077321 005</u>	Sep 09, 2005
<u>AB</u>			<u>40MG</u>	<u>A077321 006</u>	Sep 09, 2005
<u>AB</u>		PRINSTON INC	<u>2.5MG</u>	<u>A075743 001</u>	Jul 01, 2002
<u>AB</u>			<u>2.5MG</u>	<u>A076164 004</u>	Jul 01, 2002
<u>AB</u>			<u>5MG</u>	<u>A075743 002</u>	Jul 01, 2002
<u>AB</u>			<u>5MG</u>	<u>A076164 005</u>	Jul 01, 2002
<u>AB</u>			<u>10MG</u>	<u>A075743 003</u>	Jul 01, 2002
<u>AB</u>			<u>10MG</u>	<u>A076164 006</u>	Jul 01, 2002
<u>AB</u>			<u>20MG</u>	<u>A075743 004</u>	Jul 01, 2002
<u>AB</u>			<u>20MG</u>	<u>A076164 001</u>	Jul 01, 2002
<u>AB</u>			<u>30MG</u>	<u>A075743 005</u>	Jul 01, 2002
<u>AB</u>			<u>30MG</u>	<u>A076164 002</u>	Jul 01, 2002
<u>AB</u>			<u>40MG</u>	<u>A075743 006</u>	Jul 01, 2002
<u>AB</u>			<u>40MG</u>	<u>A076164 003</u>	Jul 01, 2002
<u>AB</u>		SCIEGEN PHARMS INC	<u>2.5MG</u>	<u>A212041 001</u>	Sep 15, 2020
<u>AB</u>			<u>5MG</u>	<u>A212041 002</u>	Sep 15, 2020

PRESCRIPTION DRUG PRODUCT LIST

LISINAPRIL

TABLET; ORAL

LISINAPRIL

<u>AB</u>		<u>10MG</u>	<u>A212041 003</u>	Sep 15, 2020
<u>AB</u>		<u>20MG</u>	<u>A208920 001</u>	Mar 04, 2021
<u>AB</u>		<u>30MG</u>	<u>A208920 002</u>	Mar 04, 2021
<u>AB</u>		<u>40MG</u>	<u>A208920 003</u>	Mar 04, 2021
<u>AB</u>	SUN PHARM INDS LTD	<u>2.5MG</u>	<u>A075944 001</u>	Jul 01, 2002
<u>AB</u>		<u>5MG</u>	<u>A075944 002</u>	Jul 01, 2002
<u>AB</u>		<u>10MG</u>	<u>A075944 003</u>	Jul 01, 2002
<u>AB</u>		<u>20MG</u>	<u>A075944 004</u>	Jul 01, 2002
<u>AB</u>		<u>30MG</u>	<u>A075944 006</u>	Feb 11, 2003
<u>AB</u>		<u>40MG</u>	<u>A075944 005</u>	Jul 01, 2002
<u>AB</u>	WATSON LABS	<u>2.5MG</u>	<u>A076059 001</u>	Jul 01, 2002
<u>AB</u>		<u>5MG</u>	<u>A076059 002</u>	Jul 01, 2002
<u>AB</u>		<u>10MG</u>	<u>A076059 003</u>	Jul 01, 2002
<u>AB</u>		<u>20MG</u>	<u>A076059 004</u>	Jul 01, 2002
<u>AB</u>		<u>30MG</u>	<u>A076059 005</u>	Jul 01, 2002
<u>AB</u>		<u>40MG</u>	<u>A076059 006</u>	Jul 01, 2002
<u>AB</u>	WOCKHARDT BIO AG	<u>2.5MG</u>	<u>A078402 001</u>	Apr 19, 2007
<u>AB</u>		<u>5MG</u>	<u>A078402 002</u>	Apr 19, 2007
<u>AB</u>		<u>10MG</u>	<u>A078402 003</u>	Apr 19, 2007
<u>AB</u>		<u>20MG</u>	<u>A078402 004</u>	Apr 19, 2007
<u>AB</u>		<u>30MG</u>	<u>A078402 005</u>	Apr 19, 2007
<u>AB</u>		<u>40MG</u>	<u>A078402 006</u>	Apr 19, 2007

ZESTRIL

<u>AB</u>	+	TWI PHARMS	<u>2.5MG</u>	<u>N019777 005</u>	Apr 29, 1993
<u>AB</u>	+		<u>5MG</u>	<u>N019777 001</u>	May 19, 1988
<u>AB</u>	+		<u>10MG</u>	<u>N019777 002</u>	May 19, 1988
<u>AB</u>	+		<u>20MG</u>	<u>N019777 003</u>	May 19, 1988
<u>AB</u>	+		<u>30MG</u>	<u>N019777 006</u>	Jan 20, 1999
<u>AB</u>	+		<u>40MG</u>	<u>N019777 004</u>	May 19, 1988

LITHIUM CARBONATE

CAPSULE; ORAL

LITHIUM CARBONATE

<u>AB</u>		ALEMBIC LTD	<u>150MG</u>	<u>A079159 001</u>	Jan 12, 2009
<u>AB</u>			<u>300MG</u>	<u>A079159 002</u>	Jan 12, 2009
<u>AB</u>			<u>600MG</u>	<u>A079159 003</u>	Jan 12, 2009
<u>AB</u>		GLENMARK PHARMS LTD	<u>150MG</u>	<u>A079139 001</u>	Feb 03, 2009
<u>AB</u>			<u>300MG</u>	<u>A079139 002</u>	Feb 03, 2009
<u>AB</u>			<u>600MG</u>	<u>A079139 003</u>	Feb 03, 2009
<u>AB</u>		HETERO LABS LTD III	<u>150MG</u>	<u>A090702 001</u>	Sep 25, 2009
<u>AB</u>			<u>300MG</u>	<u>A090702 002</u>	Sep 25, 2009
<u>AB</u>			<u>600MG</u>	<u>A090702 003</u>	Sep 25, 2009
<u>AB</u>	+	HIKMA	<u>150MG</u>	<u>N017812 002</u>	Jan 28, 1987
<u>AB</u>	+		<u>300MG</u>	<u>N017812 001</u>	
<u>AB</u>	+		<u>600MG</u>	<u>N017812 003</u>	Jan 28, 1987

TABLET; ORAL

LITHIUM CARBONATE

<u>AB</u>	+	HIKMA	<u>300MG</u>	<u>N018558 001</u>	Jan 29, 1982
<u>AB</u>		SUN PHARM INDS INC	<u>300MG</u>	<u>A091027 001</u>	Jun 24, 2010

TABLET, EXTENDED RELEASE; ORAL

LITHIUM CARBONATE

<u>AB</u>		GLENMARK PHARMS INC	<u>300MG</u>	<u>A091544 001</u>	Dec 27, 2010
<u>AB</u>		GLENMARK PHARMS LTD	<u>450MG</u>	<u>A091616 001</u>	Feb 14, 2011
<u>AB</u>		HERITAGE PHARMA	<u>300MG</u>	<u>A205532 001</u>	Sep 29, 2016
<u>AB</u>		HIKMA	<u>300MG</u>	<u>A076832 001</u>	Oct 28, 2004
<u>AB</u>	!		<u>450MG</u>	<u>A076691 001</u>	Jan 05, 2004
<u>AB</u>		MYLAN PHARMS INC	<u>300MG</u>	<u>A202288 001</u>	Jun 29, 2012
<u>AB</u>			<u>450MG</u>	<u>A202219 001</u>	Aug 08, 2012
<u>AB</u>		UNIQUE	<u>300MG</u>	<u>A204779 001</u>	Jul 27, 2015
<u>AB</u>			<u>450MG</u>	<u>A205663 001</u>	Jun 05, 2017

LITHOBID

<u>AB</u>	+	ANI PHARMS	<u>300MG</u>	<u>N018027 001</u>	
-----------	---	------------	--------------	--------------------	--

LITHIUM CITRATE

SYRUP; ORAL

LITHIUM CITRATE

<u>AA</u>	!	RUBICON	<u>EQ 300MG CARBONATE/5ML</u>	<u>A218036 001</u>	Aug 14, 2023
<u>AA</u>		SCIEGEN PHARMS INC	<u>EQ 300MG CARBONATE/5ML</u>	<u>A217183 001</u>	Mar 18, 2024

PRESCRIPTION DRUG PRODUCT LIST

LODOXAMIDE TROMETHAMINE

SOLUTION/DROPS;OPHTHALMIC

ALOMIDE

+! NOVARTIS

EQ 0.1% BASE

N020191 001 Sep 23, 1993

LOFEXIDINE HYDROCHLORIDE

TABLET;ORAL

LOFEXIDINE HYDROCHLORIDE**AB** INDOCOEQ 0.18MG BASEA218613 001 Aug 20, 2024LUCEMYRA**AB** +! USWMEQ 0.18MG BASEN209229 001 May 16, 2018LOMITAPIDE MESYLATE

CAPSULE;ORAL

JUXTAPID

+ CHIESI

EQ 5MG BASE

N203858 001 Dec 21, 2012

+

EQ 10MG BASE

N203858 002 Dec 21, 2012

+

EQ 20MG BASE

N203858 003 Dec 21, 2012

+

EQ 30MG BASE

N203858 004 Apr 23, 2015

LOMUSTINE

CAPSULE;ORAL

GLEOSTINE

+ LATINA PHARMA

10MG

N017588 001

+!

40MG

N017588 002

+

100MG

N017588 003

LONAFARNIB

CAPSULE;ORAL

ZOKINVY

+ SENTYNL THERAPS INC

50MG

N213969 001 Nov 20, 2020

+!

75MG

N213969 002 Nov 20, 2020

LOPERAMIDE HYDROCHLORIDE

CAPSULE;ORAL

LOPERAMIDE HYDROCHLORIDE**AB** AUROBINDO PHARMA2MGA218122 001 Sep 05, 2023

LTD

AB BIONPHARMA2MGA215579 001 Oct 22, 2021**AB** EDENBRIDGE PHARMS2MGA215001 001 Oct 06, 2021**AB** JUBILANT CADISTA2MGA217840 001 Jul 05, 2023**AB** ! MYLAN2MGA072741 001 Sep 18, 1991**AB** RUBICON2MGA216876 001 Jan 26, 2023**AB** TEVA2MGA073192 001 Apr 30, 1992**AB** ZYDUS LIFESCIENCES2MGA217471 001 Mar 23, 2023LOPINAVIR; RITONAVIR

SOLUTION;ORAL

KALETRA**AA** +! ABBVIE80MG/ML; 20MG/MLN021251 001 Sep 15, 2000LOPINAVIR AND RITONAVIR**AA** LANNETT CO INC80MG/ML; 20MG/MLA207407 001 Dec 27, 2016

TABLET;ORAL

KALETRA**AB** + ABBVIE100MG; 25MGN021906 002 Nov 09, 2007**AB** +!200MG; 50MGN021906 001 Oct 28, 2005LOPINAVIR AND RITONAVIR**AB** HETERO LABS LTD III100MG; 25MGA091677 001 Jun 04, 2021**AB**200MG; 50MGA091677 002 Jun 04, 2021**AB** LAURUS100MG; 25MGA213857 001 Mar 21, 2022**AB**200MG; 50MGA213857 002 Mar 21, 2022**AB** MACLEODS PHARMS LTD100MG; 25MGA204739 001 Jul 25, 2024**AB**200MG; 50MGA204739 002 Jul 25, 2024**AB** MYLAN LABS LTD100MG; 25MGA079074 001 Feb 07, 2024**AB**200MG; 50MGA079074 002 Feb 07, 2024LORAZEPAM

CAPSULE, EXTENDED RELEASE;ORAL

LOREEV XR

+ ALMATICA

1MG

N214826 001 Aug 27, 2021

+

1.5MG

N214826 004 Feb 16, 2022

+

2MG

N214826 002 Aug 27, 2021

+!

3MG

N214826 003 Aug 27, 2021

CONCENTRATE;ORAL

LORAZEPAM**AA** AMNEAL PHARMS2MG/MLA091383 001 Dec 23, 2009

PRESCRIPTION DRUG PRODUCT LIST

LORAZEPAM

CONCENTRATE; ORAL

LORAZEPAM

<u>AA</u>	LUPIN LTD	<u>2MG/ML</u>	<u>A091407</u>	<u>001</u>	Feb 19, 2013
<u>AA</u>	PHARM ASSOC	<u>2MG/ML</u>	<u>A090260</u>	<u>001</u>	Jun 15, 2010

LORAZEPAM INTENSOL

<u>AA</u>	! HIKMA	<u>2MG/ML</u>	<u>A072755</u>	<u>001</u>	Jun 28, 1991
-----------	---------	---------------	----------------	------------	--------------

INJECTABLE; INJECTION

ATIVAN

<u>AP</u>	+! HIKMA	<u>2MG/ML</u>	<u>N018140</u>	<u>001</u>	
<u>AP</u>	+!	<u>4MG/ML</u>	<u>N018140</u>	<u>002</u>	

LORAZEPAM

<u>AP</u>	FRESENIUS KABI USA	<u>2MG/ML</u>	<u>A217598</u>	<u>001</u>	Nov 18, 2024
<u>AP</u>	HOSPIRA	<u>2MG/ML</u>	<u>A074243</u>	<u>001</u>	Apr 12, 1994
<u>AP</u>		<u>2MG/ML</u>	<u>A074282</u>	<u>001</u>	May 27, 1994
<u>AP</u>		<u>4MG/ML</u>	<u>A074243</u>	<u>002</u>	Apr 12, 1994
<u>AP</u>		<u>4MG/ML</u>	<u>A074282</u>	<u>002</u>	May 27, 1994
<u>AP</u>	INTL MEDICATION SYS	<u>2MG/ML</u>	<u>A076150</u>	<u>001</u>	Nov 15, 2004
<u>AP</u>	RISING	<u>2MG/ML</u>	<u>A075025</u>	<u>001</u>	Jul 23, 1998

TABLET; ORAL

ATIVAN

<u>AB</u>	+ BAUSCH	<u>0.5MG</u>	<u>N017794</u>	<u>001</u>	
<u>AB</u>	+	<u>1MG</u>	<u>N017794</u>	<u>002</u>	
<u>AB</u>	+!	<u>2MG</u>	<u>N017794</u>	<u>003</u>	

LORAZEPAM

<u>AB</u>	ANI PHARMS	<u>0.5MG</u>	<u>A077396</u>	<u>001</u>	Dec 13, 2006
<u>AB</u>		<u>1MG</u>	<u>A077396</u>	<u>002</u>	Dec 13, 2006
<u>AB</u>		<u>2MG</u>	<u>A077396</u>	<u>003</u>	Dec 13, 2006
<u>AB</u>	AUROLIFE PHARMA LLC	<u>0.5MG</u>	<u>A203572</u>	<u>001</u>	Dec 22, 2017
<u>AB</u>		<u>1MG</u>	<u>A203572</u>	<u>002</u>	Dec 22, 2017
<u>AB</u>		<u>2MG</u>	<u>A203572</u>	<u>003</u>	Dec 22, 2017
<u>AB</u>	LEADING	<u>0.5MG</u>	<u>A078203</u>	<u>001</u>	Jul 30, 2007
<u>AB</u>		<u>1MG</u>	<u>A078203</u>	<u>002</u>	Jul 30, 2007
<u>AB</u>		<u>2MG</u>	<u>A078203</u>	<u>003</u>	Jul 30, 2007
<u>AB</u>	OXFORD PHARMS	<u>0.5MG</u>	<u>A077754</u>	<u>001</u>	May 10, 2006
<u>AB</u>		<u>1MG</u>	<u>A077754</u>	<u>002</u>	May 10, 2006
<u>AB</u>		<u>2MG</u>	<u>A077754</u>	<u>003</u>	May 10, 2006
<u>AB</u>	SANDOZ	<u>0.5MG</u>	<u>A071141</u>	<u>002</u>	Apr 21, 1987
<u>AB</u>		<u>1MG</u>	<u>A071141</u>	<u>003</u>	Apr 21, 1987
<u>AB</u>		<u>2MG</u>	<u>A071141</u>	<u>001</u>	Apr 21, 1987
<u>AB</u>	SUN PHARM INDS LTD	<u>0.5MG</u>	<u>A076045</u>	<u>001</u>	Aug 29, 2001
<u>AB</u>		<u>1MG</u>	<u>A076045</u>	<u>002</u>	Aug 29, 2001
<u>AB</u>		<u>2MG</u>	<u>A076045</u>	<u>003</u>	Aug 29, 2001
<u>AB</u>	WATSON LABS	<u>0.5MG</u>	<u>A072926</u>	<u>001</u>	Oct 31, 1991
<u>AB</u>		<u>1MG</u>	<u>A072927</u>	<u>001</u>	Oct 31, 1991
<u>AB</u>		<u>2MG</u>	<u>A072928</u>	<u>001</u>	Oct 31, 1991

LORLATINIB

TABLET; ORAL

LORBRENA

+	PFIZER	25MG	N210868	001	Nov 02, 2018
+!		100MG	N210868	002	Nov 02, 2018

LOSARTAN POTASSIUM

TABLET; ORAL

COZAAR

<u>AB</u>	+ ORGANON	<u>25MG</u>	<u>N020386</u>	<u>001</u>	Apr 14, 1995
<u>AB</u>	+	<u>50MG</u>	<u>N020386</u>	<u>002</u>	Apr 14, 1995
<u>AB</u>	+!	<u>100MG</u>	<u>N020386</u>	<u>003</u>	Oct 13, 1998

LOSARTAN POTASSIUM

<u>AB</u>	ALEMBIC PHARMS LTD	<u>25MG</u>	<u>A090428</u>	<u>001</u>	Oct 06, 2010
<u>AB</u>		<u>50MG</u>	<u>A090428</u>	<u>002</u>	Oct 06, 2010
<u>AB</u>		<u>100MG</u>	<u>A090428</u>	<u>003</u>	Oct 06, 2010
<u>AB</u>	APOTEX	<u>25MG</u>	<u>A218551</u>	<u>001</u>	Jun 04, 2024
<u>AB</u>		<u>50MG</u>	<u>A218551</u>	<u>002</u>	Jun 04, 2024
<u>AB</u>		<u>100MG</u>	<u>A218551</u>	<u>003</u>	Jun 04, 2024
<u>AB</u>	AUROBINDO PHARMA	<u>25MG</u>	<u>A090083</u>	<u>001</u>	Oct 06, 2010
<u>AB</u>		<u>50MG</u>	<u>A090083</u>	<u>002</u>	Oct 06, 2010
<u>AB</u>		<u>100MG</u>	<u>A090083</u>	<u>003</u>	Oct 06, 2010
<u>AB</u>	CHARTWELL RX	<u>25MG</u>	<u>A077424</u>	<u>001</u>	Oct 06, 2010
<u>AB</u>		<u>50MG</u>	<u>A077424</u>	<u>002</u>	Oct 06, 2010
<u>AB</u>		<u>100MG</u>	<u>A077424</u>	<u>003</u>	Oct 06, 2010
<u>AB</u>	GRANULES	<u>25MG</u>	<u>A215959</u>	<u>001</u>	Feb 23, 2023

PRESCRIPTION DRUG PRODUCT LIST

LOSARTAN POTASSIUM

TABLET; ORAL

LOSARTAN POTASSIUM

<u>AB</u>		<u>50MG</u>	<u>A215959 002</u>	Feb 23, 2023
<u>AB</u>		<u>100MG</u>	<u>A215959 003</u>	Feb 23, 2023
<u>AB</u>	HETERO LABS LTD V	<u>25MG</u>	<u>A203835 001</u>	Aug 12, 2015
<u>AB</u>		<u>50MG</u>	<u>A203835 002</u>	Aug 12, 2015
<u>AB</u>		<u>100MG</u>	<u>A203835 003</u>	Aug 12, 2015
<u>AB</u>	JUBILANT CADISTA	<u>25MG</u>	<u>A201170 001</u>	Sep 18, 2012
<u>AB</u>		<u>50MG</u>	<u>A201170 002</u>	Sep 18, 2012
<u>AB</u>		<u>100MG</u>	<u>A201170 003</u>	Sep 18, 2012
<u>AB</u>	LUPIN LTD	<u>25MG</u>	<u>A078232 001</u>	Oct 06, 2010
<u>AB</u>		<u>50MG</u>	<u>A078232 002</u>	Oct 06, 2010
<u>AB</u>		<u>100MG</u>	<u>A078232 003</u>	Oct 06, 2010
<u>AB</u>	MACLEODS PHARMS LTD	<u>25MG</u>	<u>A202230 001</u>	May 30, 2012
<u>AB</u>		<u>50MG</u>	<u>A202230 002</u>	May 30, 2012
<u>AB</u>		<u>100MG</u>	<u>A202230 003</u>	May 30, 2012
<u>AB</u>	MICRO LABS	<u>25MG</u>	<u>A091541 001</u>	Sep 24, 2012
<u>AB</u>		<u>50MG</u>	<u>A091541 002</u>	Sep 24, 2012
<u>AB</u>		<u>100MG</u>	<u>A091541 003</u>	Sep 24, 2012
<u>AB</u>	MSN	<u>25MG</u>	<u>A217396 001</u>	Aug 14, 2023
<u>AB</u>		<u>50MG</u>	<u>A217396 002</u>	Aug 14, 2023
<u>AB</u>		<u>100MG</u>	<u>A217396 003</u>	Aug 14, 2023
<u>AB</u>	PRINSTON INC	<u>25MG</u>	<u>A091497 001</u>	Jun 06, 2011
<u>AB</u>		<u>50MG</u>	<u>A091497 002</u>	Jun 06, 2011
<u>AB</u>		<u>100MG</u>	<u>A091497 003</u>	Jun 06, 2011
<u>AB</u>	STRIDES PHARMA	<u>25MG</u>	<u>A090382 001</u>	Oct 06, 2010
<u>AB</u>		<u>50MG</u>	<u>A090382 002</u>	Oct 06, 2010
<u>AB</u>		<u>100MG</u>	<u>A090382 003</u>	Oct 06, 2010
<u>AB</u>	UNICHEM	<u>25MG</u>	<u>A203030 001</u>	Oct 14, 2015
<u>AB</u>		<u>50MG</u>	<u>A203030 002</u>	Oct 14, 2015
<u>AB</u>		<u>100MG</u>	<u>A203030 003</u>	Oct 14, 2015
<u>AB</u>	WATSON LABS	<u>25MG</u>	<u>A091129 001</u>	Oct 06, 2010
<u>AB</u>		<u>50MG</u>	<u>A091129 002</u>	Oct 06, 2010
<u>AB</u>		<u>100MG</u>	<u>A091129 003</u>	Oct 06, 2010
<u>AB</u>	ZYDUS PHARMS USA INC	<u>25MG</u>	<u>A078243 001</u>	Oct 06, 2010
<u>AB</u>		<u>50MG</u>	<u>A078243 002</u>	Oct 06, 2010
<u>AB</u>		<u>100MG</u>	<u>A078243 003</u>	Oct 06, 2010

LOTEPREDNOL ETABONATE

GEL; OPHTHALMIC

LOTEMAX

<u>AB</u>	<u>+</u> !	BAUSCH AND LOMB INC	<u>0.5%</u>	<u>N202872 001</u>	Sep 28, 2012
-----------	------------	---------------------	-------------	--------------------	--------------

LOTEPREDNOL ETABONATE

<u>AB</u>		SENTISS	<u>0.5%</u>	<u>A212080 001</u>	Feb 10, 2021
<u>AB</u>		SUN PHARM	<u>0.5%</u>	<u>A215384 001</u>	Aug 09, 2023

LOTEMAX SM

	<u>+</u> !	BAUSCH AND LOMB INC	0.38%	N208219 001	Feb 22, 2019
--	------------	---------------------	-------	-------------	--------------

OINTMENT; OPHTHALMIC

LOTEMAX

	<u>+</u> !	BAUSCH AND LOMB	0.5%	N200738 001	Apr 15, 2011
--	------------	-----------------	------	-------------	--------------

SUSPENSION/DROPS; OPHTHALMIC

ALREX

<u>AB</u>	<u>+</u> !	BAUSCH AND LOMB	<u>0.2%</u>	<u>N020803 001</u>	Mar 09, 1998
-----------	------------	-----------------	-------------	--------------------	--------------

LOTEMAX

<u>AB</u>	<u>+</u> !	BAUSCH AND LOMB	<u>0.5%</u>	<u>N020583 001</u>	Mar 09, 1998
-----------	------------	-----------------	-------------	--------------------	--------------

LOTEPREDNOL ETABONATE

<u>AB</u>		AMNEAL	<u>0.2%</u>	<u>A216345 001</u>	Dec 13, 2024
<u>AB</u>		LUPIN LTD	<u>0.2%</u>	<u>A215550 001</u>	Dec 26, 2023
<u>AB</u>			<u>0.5%</u>	<u>A215719 001</u>	Apr 24, 2024
<u>AB</u>		PADAGIS US	<u>0.2%</u>	<u>A215203 001</u>	Jul 10, 2024
<u>AB</u>		SENTISS	<u>0.2%</u>	<u>A215933 001</u>	Apr 12, 2023
<u>AB</u>			<u>0.5%</u>	<u>A207609 001</u>	Apr 17, 2019
<u>AB</u>		SUN PHARM	<u>0.5%</u>	<u>A212450 001</u>	Feb 26, 2021
		EYSUVIS			
	<u>+</u> !	ALCON LABS INC	0.25%	N210933 001	Oct 26, 2020
		INVELTYS			
	<u>+</u> !	ALCON LABS INC	1%	N210565 001	Aug 22, 2018

PRESCRIPTION DRUG PRODUCT LIST

LOTEPREDNOL ETABONATE; TOBRAMYCIN

SUSPENSION/DROPS;OPHTHALMIC

ZYLET

+! BAUSCH AND LOMB 0.5%;0.3% N050804 001 Dec 14, 2004

LOTILANER

SOLUTION/DROPS;OPHTHALMIC

XDEMVI

+! TARSUS 0.25% N217603 001 Jul 24, 2023

LOVASTATIN

TABLET;ORAL

LOVASTATIN

<u>AB</u>	ACTAVIS ELIZABETH	<u>10MG</u>	<u>A075828 001</u>	Dec 17, 2001
<u>AB</u>		<u>20MG</u>	<u>A075828 002</u>	Dec 17, 2001
<u>AB</u>		<u>40MG</u>	<u>A075828 003</u>	Dec 17, 2001
<u>AB</u>	CARLSBAD	<u>10MG</u>	<u>A075991 001</u>	Jun 05, 2002
<u>AB</u>		<u>20MG</u>	<u>A075991 002</u>	Jun 05, 2002
<u>AB</u>	!	<u>40MG</u>	<u>A075991 003</u>	Jun 05, 2002
<u>AB</u>	CHARTWELL RX	<u>10MG</u>	<u>A075300 001</u>	Dec 17, 2001
<u>AB</u>		<u>20MG</u>	<u>A075300 002</u>	Dec 17, 2001
<u>AB</u>		<u>40MG</u>	<u>A075300 003</u>	Dec 17, 2001
<u>AB</u>	COREPHARMA	<u>10MG</u>	<u>A077748 001</u>	Feb 28, 2007
<u>AB</u>		<u>20MG</u>	<u>A077748 002</u>	Feb 28, 2007
<u>AB</u>		<u>40MG</u>	<u>A077748 003</u>	Feb 28, 2007
<u>AB</u>	EPIC PHARMA LLC	<u>10MG</u>	<u>A075636 001</u>	Dec 17, 2001
<u>AB</u>		<u>20MG</u>	<u>A075636 002</u>	Dec 17, 2001
<u>AB</u>		<u>40MG</u>	<u>A075636 003</u>	Dec 17, 2001
<u>AB</u>	LUPIN	<u>10MG</u>	<u>A078296 001</u>	Mar 14, 2008
<u>AB</u>		<u>20MG</u>	<u>A078296 002</u>	Nov 01, 2007
<u>AB</u>		<u>40MG</u>	<u>A078296 003</u>	Nov 01, 2007
<u>AB</u>	TEVA	<u>10MG</u>	<u>A075551 003</u>	Dec 17, 2001
<u>AB</u>		<u>20MG</u>	<u>A075551 002</u>	Dec 17, 2001
<u>AB</u>		<u>40MG</u>	<u>A075551 001</u>	Dec 17, 2001

TABLET, EXTENDED RELEASE;ORAL

ALTOPREV

+	COVIS	20MG	N021316 002	Jun 26, 2002
+		40MG	N021316 003	Jun 26, 2002
+!		60MG	N021316 004	Jun 26, 2002

LOXAPINE

POWDER; INHALATION

ADASUVE

+! ALEXZA PHARMS 10MG N022549 001 Dec 21, 2012

LOXAPINE SUCCINATE

CAPSULE;ORAL

LOXAPINE SUCCINATE

<u>AB</u>	ELITE LABS INC	<u>EQ 5MG BASE</u>	<u>A076868 001</u>	Aug 04, 2005
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A076868 002</u>	Aug 04, 2005
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>A076868 003</u>	Aug 04, 2005
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A076868 004</u>	Aug 04, 2005
<u>AB</u>	LANNETT CO INC	<u>EQ 5MG BASE</u>	<u>A090695 001</u>	Sep 26, 2011
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A090695 002</u>	Sep 26, 2011
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>A090695 003</u>	Sep 26, 2011
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A090695 004</u>	Sep 26, 2011
<u>AB</u>	WATSON LABS	<u>EQ 5MG BASE</u>	<u>A072204 001</u>	Jun 15, 1988
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A072205 001</u>	Jun 15, 1988
<u>AB</u>	!	<u>EQ 25MG BASE</u>	<u>A072206 001</u>	Jun 15, 1988
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A072062 001</u>	Jun 15, 1988

LUBIPROSTONE

CAPSULE;ORAL

AMITIZA

<u>AB</u>	+	SUCAMPO PHARMA LLC	<u>8MCG</u>	<u>N021908 002</u>	Apr 29, 2008
<u>AB</u>	+!		<u>24MCG</u>	<u>N021908 001</u>	Jan 31, 2006

LUBIPROSTONE

<u>AB</u>	AMNEAL	<u>8MCG</u>	<u>A209450 001</u>	Nov 30, 2021
<u>AB</u>		<u>24MCG</u>	<u>A209450 002</u>	Nov 30, 2021
<u>AB</u>	ASCENT PHARMS INC	<u>8MCG</u>	<u>A218640 001</u>	Jan 02, 2025
<u>AB</u>		<u>24MCG</u>	<u>A218640 002</u>	Jan 02, 2025
<u>AB</u>	DR REDDYS	<u>8MCG</u>	<u>A206994 001</u>	Feb 08, 2022
<u>AB</u>		<u>24MCG</u>	<u>A206994 002</u>	Feb 08, 2022
<u>AB</u>	TEVA PHARMS USA INC	<u>8MCG</u>	<u>A209920 001</u>	Jan 18, 2022
<u>AB</u>		<u>24MCG</u>	<u>A209920 002</u>	Jan 18, 2022

PRESCRIPTION DRUG PRODUCT LIST

LUBIPROSTONE

CAPSULE; ORAL

LUBIPROSTONE

<u>AB</u>	ZYDUS PHARMS	<u>8MCG</u>	<u>A214131 001</u>	Mar 23, 2023
<u>AB</u>		<u>24MCG</u>	<u>A214131 002</u>	Mar 23, 2023

LULICONAZOLE

CREAM; TOPICAL

LUZU

+	BAUSCH	1%	N204153 001	Nov 14, 2013
---	--------	----	-------------	--------------

LUMASIRAN SODIUM

SOLUTION; SUBCUTANEOUS

OXLUMO

+	ALNYLAM PHARMS INC	EQ 94.5MG BASE/0.5ML (EQ 94.5MG BASE/0.5ML)	N214103 001	Nov 23, 2020
---	--------------------	---	-------------	--------------

LUMATEPERONE TOSYLATE

CAPSULE; ORAL

CAPLYTA

+	INTRA-CELLULAR	EQ 10.5MG BASE	N209500 002	Apr 22, 2022
+		EQ 21MG BASE	N209500 003	Apr 22, 2022
+		EQ 42MG BASE	N209500 001	Dec 20, 2019

LURASIDONE HYDROCHLORIDE

TABLET; ORAL

LATUDA

<u>AB</u>	+	SUNOVION PHARMS INC	<u>20MG</u>	<u>N200603 003</u>	Dec 07, 2011
<u>AB</u>	+		<u>40MG</u>	<u>N200603 001</u>	Oct 28, 2010
<u>AB</u>	+		<u>60MG</u>	<u>N200603 005</u>	Jul 12, 2013
<u>AB</u>	+		<u>80MG</u>	<u>N200603 002</u>	Oct 28, 2010
<u>AB</u>	+		<u>120MG</u>	<u>N200603 004</u>	Apr 26, 2012

LURASIDONE HYDROCHLORIDE

<u>AB</u>		ACCORD HLTHCARE	<u>20MG</u>	<u>A208049 001</u>	Jan 03, 2019
<u>AB</u>			<u>40MG</u>	<u>A208049 002</u>	Jan 03, 2019
<u>AB</u>			<u>60MG</u>	<u>A208049 003</u>	Jan 03, 2019
<u>AB</u>			<u>80MG</u>	<u>A208049 004</u>	Jan 03, 2019
<u>AB</u>			<u>120MG</u>	<u>A208049 005</u>	Jan 03, 2019
<u>AB</u>		ADAPTIS	<u>20MG</u>	<u>A212091 001</u>	Dec 28, 2020
<u>AB</u>			<u>40MG</u>	<u>A212091 002</u>	Dec 28, 2020
<u>AB</u>			<u>60MG</u>	<u>A212091 003</u>	Dec 28, 2020
<u>AB</u>			<u>80MG</u>	<u>A212091 004</u>	Dec 28, 2020
<u>AB</u>			<u>120MG</u>	<u>A212091 005</u>	Dec 28, 2020
<u>AB</u>		ALEMBIC	<u>20MG</u>	<u>A213248 001</u>	May 13, 2021
<u>AB</u>			<u>40MG</u>	<u>A213248 002</u>	May 13, 2021
<u>AB</u>			<u>60MG</u>	<u>A213248 003</u>	May 13, 2021
<u>AB</u>			<u>80MG</u>	<u>A213248 004</u>	May 13, 2021
<u>AB</u>			<u>120MG</u>	<u>A213248 005</u>	May 13, 2021
<u>AB</u>		ALKEM LABS LTD	<u>20MG</u>	<u>A212244 001</u>	Dec 13, 2022
<u>AB</u>			<u>40MG</u>	<u>A212244 002</u>	Dec 13, 2022
<u>AB</u>			<u>60MG</u>	<u>A212244 003</u>	Dec 13, 2022
<u>AB</u>			<u>80MG</u>	<u>A212244 004</u>	Dec 13, 2022
<u>AB</u>			<u>120MG</u>	<u>A212244 005</u>	Dec 13, 2022
<u>AB</u>		AMNEAL PHARMS CO	<u>20MG</u>	<u>A208002 001</u>	Jan 03, 2019
<u>AB</u>			<u>40MG</u>	<u>A208002 002</u>	Jan 03, 2019
<u>AB</u>			<u>60MG</u>	<u>A208002 003</u>	Jan 03, 2019
<u>AB</u>			<u>80MG</u>	<u>A208002 004</u>	Jan 03, 2019
<u>AB</u>			<u>120MG</u>	<u>A208002 005</u>	Jan 03, 2019
<u>AB</u>		ANNORA PHARMA	<u>20MG</u>	<u>A218174 001</u>	Aug 21, 2024
<u>AB</u>			<u>40MG</u>	<u>A218174 002</u>	Aug 21, 2024
<u>AB</u>			<u>60MG</u>	<u>A218174 003</u>	Aug 21, 2024
<u>AB</u>			<u>80MG</u>	<u>A218174 004</u>	Aug 21, 2024
<u>AB</u>			<u>120MG</u>	<u>A218174 005</u>	Aug 21, 2024
<u>AB</u>		AUROBINDO PHARMA LTD	<u>20MG</u>	<u>A208045 001</u>	Mar 10, 2023
<u>AB</u>			<u>40MG</u>	<u>A208045 002</u>	Mar 10, 2023
<u>AB</u>			<u>60MG</u>	<u>A208045 003</u>	Mar 10, 2023
<u>AB</u>			<u>80MG</u>	<u>A208045 004</u>	Mar 10, 2023
<u>AB</u>			<u>120MG</u>	<u>A208045 005</u>	Mar 10, 2023
<u>AB</u>		DR REDDYS	<u>20MG</u>	<u>A208047 001</u>	Aug 24, 2021
<u>AB</u>			<u>40MG</u>	<u>A208047 002</u>	Aug 24, 2021
<u>AB</u>			<u>60MG</u>	<u>A208047 003</u>	Aug 24, 2021
<u>AB</u>			<u>80MG</u>	<u>A208047 004</u>	Aug 24, 2021
<u>AB</u>			<u>20MG</u>	<u>A208058 001</u>	Sep 04, 2019

PRESCRIPTION DRUG PRODUCT LIST

LURASIDONE HYDROCHLORIDE

TABLET; ORAL

LURASIDONE HYDROCHLORIDE

<u>AB</u>		<u>40MG</u>	<u>A208058</u>	<u>002</u>	Sep 04, 2019
<u>AB</u>		<u>60MG</u>	<u>A208058</u>	<u>003</u>	Sep 04, 2019
<u>AB</u>		<u>80MG</u>	<u>A208058</u>	<u>004</u>	Sep 04, 2019
<u>AB</u>		<u>120MG</u>	<u>A208058</u>	<u>005</u>	Feb 28, 2023
<u>AB</u>	INVAGEN PHARMS	<u>20MG</u>	<u>A208028</u>	<u>001</u>	Jan 03, 2019
<u>AB</u>		<u>40MG</u>	<u>A208028</u>	<u>002</u>	Jan 03, 2019
<u>AB</u>		<u>60MG</u>	<u>A208028</u>	<u>003</u>	Jan 03, 2019
<u>AB</u>		<u>80MG</u>	<u>A208028</u>	<u>004</u>	Jan 03, 2019
<u>AB</u>		<u>120MG</u>	<u>A208028</u>	<u>005</u>	Jan 03, 2019
<u>AB</u>	JUBILANT GENERICS	<u>20MG</u>	<u>A210388</u>	<u>001</u>	Dec 06, 2024
<u>AB</u>		<u>40MG</u>	<u>A210388</u>	<u>002</u>	Dec 06, 2024
<u>AB</u>		<u>60MG</u>	<u>A210388</u>	<u>003</u>	Dec 06, 2024
<u>AB</u>		<u>80MG</u>	<u>A210388</u>	<u>004</u>	Dec 06, 2024
<u>AB</u>		<u>120MG</u>	<u>A210388</u>	<u>005</u>	Dec 06, 2024
<u>AB</u>	LUPIN LTD	<u>20MG</u>	<u>A208031</u>	<u>001</u>	Jan 03, 2019
<u>AB</u>		<u>40MG</u>	<u>A208031</u>	<u>002</u>	Jan 03, 2019
<u>AB</u>		<u>60MG</u>	<u>A208031</u>	<u>003</u>	Jan 03, 2019
<u>AB</u>		<u>80MG</u>	<u>A208031</u>	<u>004</u>	Jan 03, 2019
<u>AB</u>		<u>120MG</u>	<u>A208031</u>	<u>005</u>	Jan 03, 2019
<u>AB</u>	MACLEODS PHARMS LTD	<u>20MG</u>	<u>A212124</u>	<u>001</u>	Jun 09, 2023
<u>AB</u>		<u>40MG</u>	<u>A212124</u>	<u>002</u>	Jun 09, 2023
<u>AB</u>		<u>60MG</u>	<u>A212124</u>	<u>003</u>	Jun 09, 2023
<u>AB</u>		<u>80MG</u>	<u>A212124</u>	<u>004</u>	Jun 09, 2023
<u>AB</u>		<u>120MG</u>	<u>A212124</u>	<u>005</u>	Jun 09, 2023
<u>AB</u>	MSN	<u>20MG</u>	<u>A208037</u>	<u>001</u>	Sep 09, 2022
<u>AB</u>		<u>40MG</u>	<u>A208037</u>	<u>002</u>	Sep 09, 2022
<u>AB</u>		<u>60MG</u>	<u>A208037</u>	<u>003</u>	Sep 09, 2022
<u>AB</u>		<u>80MG</u>	<u>A208037</u>	<u>004</u>	Sep 09, 2022
<u>AB</u>		<u>120MG</u>	<u>A208037</u>	<u>005</u>	Sep 09, 2022
<u>AB</u>	SUN PHARM	<u>20MG</u>	<u>A208066</u>	<u>001</u>	Jan 04, 2019
<u>AB</u>		<u>40MG</u>	<u>A208066</u>	<u>002</u>	Jan 04, 2019
<u>AB</u>		<u>60MG</u>	<u>A208066</u>	<u>003</u>	Jan 04, 2019
<u>AB</u>		<u>80MG</u>	<u>A208066</u>	<u>004</u>	Jan 04, 2019
<u>AB</u>		<u>120MG</u>	<u>A208066</u>	<u>005</u>	Jan 04, 2019
<u>AB</u>	TORRENT	<u>20MG</u>	<u>A208055</u>	<u>001</u>	Jan 03, 2019
<u>AB</u>		<u>40MG</u>	<u>A208055</u>	<u>002</u>	Jan 03, 2019
<u>AB</u>		<u>80MG</u>	<u>A208055</u>	<u>003</u>	Jan 03, 2019
<u>AB</u>		<u>120MG</u>	<u>A208055</u>	<u>004</u>	Jan 03, 2019
<u>AB</u>	ZYDUS PHARMS	<u>20MG</u>	<u>A208052</u>	<u>001</u>	Mar 19, 2019
<u>AB</u>		<u>40MG</u>	<u>A208052</u>	<u>002</u>	Mar 19, 2019
<u>AB</u>		<u>60MG</u>	<u>A208052</u>	<u>003</u>	Mar 19, 2019
<u>AB</u>		<u>80MG</u>	<u>A208052</u>	<u>004</u>	Mar 19, 2019
<u>AB</u>		<u>120MG</u>	<u>A208052</u>	<u>005</u>	Mar 19, 2019

LURBINECTEDIN

POWDER; INTRAVENOUS

ZEPZELCA

+! JAZZ

4MG/VIAL

N213702 001 Jun 15, 2020

LUSUTROMBOPAG

TABLET; ORAL

MULPLETA

+! VANCOCIN ITALIA

3MG

N210923 001 Jul 31, 2018

LUTETIUM LU 177 DOTATATE

SOLUTION; INTRAVENOUS

LUTATHERA

+! AAA USA INC

10mCi/ML

N208700 001 Jan 26, 2018

LUTETIUM LU-177 VIPIVOTIDE TETRAJETAN

SOLUTION; INTRAVENOUS

PLUVICTO

+! NOVARTIS

27mCi/ML

N215833 001 Mar 23, 2022

MACITENTAN

TABLET; ORAL

OPSUMIT

+! ACTELION

10MG

N204410 001 Oct 18, 2013

PRESCRIPTION DRUG PRODUCT LIST

MACITENTAN; TADALAFIL

TABLET; ORAL

OPSYNVI

+ ACTELION

10MG; 20MG

N218490 001 Mar 22, 2024

+!

10MG; 40MG

N218490 002 Mar 22, 2024

MAFENIDE ACETATE

CREAM; TOPICAL

SULFAMYLON

+! RISING

EQ 85MG BASE/GM

N016763 001

MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; POTASSIUM PHOSPHATE, MONOBASIC; SODIUM ACETATE; SODIUM CHLORIDE; SODIUM GLUCONATE; SODIUM PHOSPHATE, DIBASIC, HEPTAHYDRATE

INJECTABLE; INJECTION

ISOLYTE S PH 7.4 IN PLASTIC CONTAINER

+! B BRAUN

30MG/100ML; 37MG/100ML; 0.82MG/100ML; 370MG/100ML; 530MG/100ML; 500MG/100ML; 12MG/100ML

N019696 001 Sep 29, 1989

MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE; SODIUM GLUCONATE

INJECTABLE; INJECTION

MULTIPLE ELECTROLYTES INJECTION TYPE 1 USP PH 5.5**AP** FRESENIUS KABI USA**30MG/100ML; 37MG/100ML; 368MG/100ML; 526MG/100ML; 502MG/100ML****A215370 001** Jun 29, 2022MULTIPLE ELECTROLYTES INJECTION TYPE 1 USP PH 7.4**AP** FRESENIUS KABI USA**30MG/100ML; 37MG/100ML; 368MG/100ML; 526MG/100ML; 502MG/100ML****A215371 001** Jun 08, 2022PLASMA-LYTE 148 IN WATER IN PLASTIC CONTAINER**AP** +! BAXTER HLTHCARE**30MG/100ML; 37MG/100ML; 368MG/100ML; 526MG/100ML; 502MG/100ML****N017378 001**PLASMA-LYTE A IN PLASTIC CONTAINER**AP** +! BAXTER HLTHCARE**30MG/100ML; 37MG/100ML; 368MG/100ML; 526MG/100ML; 502MG/100ML****N017378 002** Nov 22, 1982

ISOLYTE S IN PLASTIC CONTAINER

B BRAUN

30MG/100ML; 37MG/100ML; 370MG/100ML; 530MG/100ML; 500MG/100ML

N019711 001 Sep 29, 1989

NORMOSOL-R IN PLASTIC CONTAINER

ICU MEDICAL INC

30MG/100ML; 37MG/100ML; 222MG/100ML; 526MG/100ML; 502MG/100ML

N017586 001

SOLUTION; IRRIGATION

PHYSIOLYTE IN PLASTIC CONTAINER

B BRAUN

30MG/100ML; 37MG/100ML; 370MG/100ML; 530MG/100ML; 500MG/100ML

N019024 001 Jun 08, 1984

PHYSIOSOL IN PLASTIC CONTAINER

ICU MEDICAL INC

30MG/100ML; 37MG/100ML; 222MG/100ML; 526MG/100ML; 502MG/100ML

N017637 002 Jul 08, 1982

MAGNESIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE

SOLUTION; INJECTION

NORMOCARB HF 25

+! DIALYSIS SUPS

0.21GM/100ML; 2.8GM/100ML; 9.07GM/100ML

N021910 001 Jul 26, 2006

NORMOCARB HF 35

+! DIALYSIS SUPS

0.21GM/100ML; 3.97GM/100ML; 8.3GM/100ML

N021910 002 Jul 26, 2006

MAGNESIUM SULFATE

INJECTABLE; INJECTION

MAGNESIUM SULFATE**AP** B BRAUN MEDICAL INC**2GM/50ML (40MG/ML)****A207967 001** Apr 26, 2021**AP****4GM/100ML (40MG/ML)****A207967 002** Apr 26, 2021**AP****4GM/50ML (80MG/ML)****A207967 003** Apr 26, 2021MAGNESIUM SULFATE IN DEXTROSE 5% IN PLASTIC CONTAINER**AP** B BRAUN MEDICAL INC**1GM/100ML****A207966 001** Oct 27, 2020**AP** BAXTER HLTHCARE**1GM/100ML****A211965 001** Aug 11, 2020

CORP

AP FRESENIUS KABI USA**1GM/100ML****A206486 001** Mar 07, 2016**AP** +! HOSPIRA**1GM/100ML****N020488 001** Jul 11, 1995**AP** HQ SPCLT PHARMA**1GM/100ML****A207349 001** Mar 02, 2016**AP** MYLAN LABS LTD**1GM/100ML****A209932 001** Sep 10, 2018MAGNESIUM SULFATE IN PLASTIC CONTAINER**AP** AMNEAL**2GM/50ML (40MG/ML)****A216597 001** Feb 15, 2023**AP****4GM/100ML (40MG/ML)****A216597 002** Feb 15, 2023**AP** BAXTER HLTHCARE**2GM/50ML (40MG/ML)****A211966 001** Jun 01, 2020

CORP

AP**4GM/50ML (80MG/ML)****A211966 002** Jun 01, 2020**AP****4GM/100ML (40MG/ML)****A211966 003** Jun 01, 2020**AP** FRESENIUS KABI USA**4GM/100ML (40MG/ML)****A206485 001** Mar 15, 2016**AP****4GM/50ML (80MG/ML)****A206485 002** Mar 15, 2016

PRESCRIPTION DRUG PRODUCT LISTMAGNESIUM SULFATE

INJECTABLE; INJECTION

MAGNESIUM SULFATE IN PLASTIC CONTAINER

<u>AP</u>		<u>2GM/50ML (40MG/ML)</u>	<u>A206485 003</u>	Mar 15, 2016
<u>AP</u>		<u>20GM/500ML (40MG/ML)</u>	<u>A206485 004</u>	Mar 15, 2016
<u>AP</u>		<u>40GM/1000ML (40MG/ML)</u>	<u>A206485 005</u>	Mar 15, 2016
<u>AP</u>	GLAND PHARMA LTD	<u>2GM/50ML (40MG/ML)</u>	<u>A213917 001</u>	Jul 10, 2020
<u>AP</u>		<u>4GM/100ML (40MG/ML)</u>	<u>A213917 002</u>	Jul 10, 2020
<u>AP</u>	+ HOSPIRA	<u>2GM/50ML (40MG/ML)</u>	<u>N020309 003</u>	Jan 26, 2007
<u>AP</u>	+!	<u>4GM/100ML (40MG/ML)</u>	<u>N020309 001</u>	Jun 24, 1994
<u>AP</u>	+!	<u>4GM/50ML (80MG/ML)</u>	<u>N020309 002</u>	Jun 24, 1994
<u>AP</u>	+	<u>20GM/500ML (40MG/ML)</u>	<u>N020309 004</u>	Jan 18, 1995
<u>AP</u>	+	<u>40GM/1000ML (40MG/ML)</u>	<u>N020309 005</u>	Jan 18, 1995
<u>AP</u>	HQ SPCLT PHARMA	<u>2GM/50ML (40MG/ML)</u>	<u>A207350 001</u>	Dec 06, 2017
<u>AP</u>		<u>4GM/100ML (40MG/ML)</u>	<u>A207350 002</u>	Dec 06, 2017
<u>AP</u>		<u>4GM/50ML (80MG/ML)</u>	<u>A207350 003</u>	Dec 06, 2017
<u>AP</u>		<u>20GM/500ML (40MG/ML)</u>	<u>A207350 004</u>	Dec 06, 2017
<u>AP</u>		<u>40GM/1000ML (40MG/ML)</u>	<u>A207350 005</u>	Dec 06, 2017
<u>AP</u>	MILLA PHARMS	<u>2GM/50ML (40MG/ML)</u>	<u>A209642 001</u>	Nov 08, 2021
<u>AP</u>		<u>4GM/100ML (40MG/ML)</u>	<u>A209642 002</u>	Nov 08, 2021
<u>AP</u>		<u>4GM/50ML (80MG/ML)</u>	<u>A209642 003</u>	Nov 08, 2021
<u>AP</u>	MYLAN LABS LTD	<u>2GM/50ML (40MG/ML)</u>	<u>A209911 001</u>	Sep 14, 2018
<u>AP</u>		<u>4GM/100ML (40MG/ML)</u>	<u>A209911 002</u>	Sep 14, 2018

MAGNESIUM SULFATE IN DEXTROSE 5% IN PLASTIC CONTAINER

+! HOSPIRA 2GM/100ML N020488 002 Jul 11, 1995

SOLUTION; INTRAMUSCULAR, INTRAVENOUS

MAGNESIUM SULFATE

<u>AP</u>	+!	FRESENIUS KABI USA	<u>5GM/10ML (500MG/ML)</u>	<u>N019316 001</u>	Sep 08, 1986
<u>AP</u>	+!		<u>10GM/20ML (500MG/ML)</u>	<u>N019316 003</u>	Jan 29, 2016
<u>AP</u>	!	HOSPIRA	<u>5GM/10ML (500MG/ML)</u>	<u>A075151 001</u>	Apr 25, 2000
<u>AP</u>			<u>10GM/20ML (500MG/ML)</u>	<u>A202411 001</u>	May 14, 2015
	+!	FRESENIUS KABI USA	1GM/2ML (500MG/ML)	N019316 002	Sep 08, 1986
	+!		25GM/50ML (500MG/ML)	N019316 004	Jan 29, 2016

MAGNESIUM SULFATE; POLYETHYLENE GLYCOL 3350; POTASSIUM CHLORIDE; SODIUM CHLORIDE; SODIUM SULFATE

FOR SOLUTION; ORAL

SUFLAVE

+! BRAINTREE LABS 0.9GM/BOT; 178.7GM/BOT; 1.12GM/BOT; 0.5GM/BOT; 7.3GM/BOT N215344 001 Jun 15, 2023

MAGNESIUM SULFATE; POTASSIUM CHLORIDE; POTASSIUM PHOSPHATE, MONOBASIC; SODIUM CHLORIDE; SODIUM PHOSPHATE

SOLUTION; IRRIGATION

TIS-U-SOL

<u>AT</u>	BAXTER HLTHCARE	<u>20MG/100ML; 40MG/100ML; 6.25MG/100ML; 800MG/100ML; 8.75MG/100ML</u>	<u>N018508 001</u>	Feb 19, 1982
-----------	-----------------	--	--------------------	--------------

TIS-U-SOL IN PLASTIC CONTAINER

<u>AT</u>	BAXTER HLTHCARE	<u>20MG/100ML; 40MG/100ML; 6.25MG/100ML; 800MG/100ML; 8.75MG/100ML</u>	<u>N018336 001</u>	
-----------	-----------------	--	--------------------	--

MAGNESIUM SULFATE; POTASSIUM CHLORIDE; SODIUM SULFATE

TABLET; ORAL

SUTAB

+! BRAINTREE LABS 0.225GM; 0.188GM; 1.479GM N213135 001 Nov 10, 2020

MAGNESIUM SULFATE; POTASSIUM SULFATE; SODIUM SULFATE

SOLUTION; ORAL

SODIUM SULFATE, POTASSIUM SULFATE AND MAGNESIUM SULFATE

<u>AA</u>	ALKEM LABS LTD	<u>1.6GM/BOT; 3.13GM/BOT; 17.5GM/BOT</u>	<u>A213924 001</u>	Dec 29, 2023	
<u>AA</u>	ANNORA PHARMA	<u>1.6GM/BOT; 3.13GM/BOT; 17.5GM/BOT</u>	<u>A212903 001</u>	Oct 18, 2024	
<u>AA</u>	BIONPHARMA	<u>1.6GM/BOT; 3.13GM/BOT; 17.5GM/BOT</u>	<u>A204135 001</u>	Jun 24, 2024	
<u>AA</u>	NOVEL LABS INC	<u>1.6GM/BOT; 3.13GM/BOT; 17.5GM/BOT</u>	<u>A202511 001</u>	Feb 23, 2017	
<u>AA</u>	PAI HOLDINGS	<u>1.6GM/BOT; 3.13GM/BOT; 17.5GM/BOT</u>	<u>A209941 001</u>	Jul 15, 2024	
<u>AA</u>	STRIDES PHARMA	<u>1.6GM/BOT; 3.13GM/BOT; 17.5GM/BOT</u>	<u>A215469 001</u>	Nov 22, 2023	
<u>AA</u>	TARO	<u>1.6GM/BOT; 3.13GM/BOT; 17.5GM/BOT</u>	<u>A206431 001</u>	Mar 19, 2024	
<u>AA</u>	+!	BRAINTREE LABS	<u>1.6GM/BOT; 3.13GM/BOT; 17.5GM/BOT</u>	<u>N022372 001</u>	Aug 05, 2010

MALATHION

LOTION; TOPICAL

MALATHION

! SUVEN PHARMS 0.5% A091559 001 May 23, 2012

PRESCRIPTION DRUG PRODUCT LIST

MANGANESE CHLORIDE

INJECTABLE; INJECTION

MANGANESE CHLORIDE IN PLASTIC CONTAINER

+	!	HOSPIRA	EQ 0.1MG MANGANESE/ML	N018962	001	Jun 26, 1986
---	---	---------	-----------------------	---------	-----	--------------

MANNITOL

INJECTABLE; INJECTION

MANNITOL 10% IN PLASTIC CONTAINER

<u>AP</u>		B BRAUN	<u>10GM/100ML</u>	<u>N020006</u>	<u>002</u>	Jul 26, 1993
-----------	--	---------	-------------------	----------------	------------	--------------

MANNITOL 15% IN PLASTIC CONTAINER

<u>AP</u>		B BRAUN	<u>15GM/100ML</u>	<u>N020006</u>	<u>003</u>	Jul 26, 1993
-----------	--	---------	-------------------	----------------	------------	--------------

MANNITOL 20% IN PLASTIC CONTAINER

<u>AP</u>		B BRAUN	<u>20GM/100ML</u>	<u>N020006</u>	<u>004</u>	Jul 26, 1993
-----------	--	---------	-------------------	----------------	------------	--------------

<u>AP</u>		ICU MEDICAL INC	<u>20GM/100ML</u>	<u>N019603</u>	<u>004</u>	Jan 08, 1990
-----------	--	-----------------	-------------------	----------------	------------	--------------

MANNITOL 25%

<u>AP</u>		FRESENIUS KABI USA	<u>12.5GM/50ML</u>	<u>A080677</u>	<u>001</u>	
-----------	--	--------------------	--------------------	----------------	------------	--

<u>AP</u>		HOSPIRA	<u>12.5GM/50ML</u>	<u>N016269</u>	<u>006</u>	Aug 25, 1994
-----------	--	---------	--------------------	----------------	------------	--------------

MANNITOL 5% IN PLASTIC CONTAINER

<u>AP</u>		B BRAUN	<u>5GM/100ML</u>	<u>N020006</u>	<u>001</u>	Jul 26, 1993
-----------	--	---------	------------------	----------------	------------	--------------

OSMITROL 10% IN WATER

<u>AP</u>		BAXTER HLTHCARE	<u>10GM/100ML</u>	<u>N013684</u>	<u>002</u>	
-----------	--	-----------------	-------------------	----------------	------------	--

OSMITROL 10% IN WATER IN PLASTIC CONTAINER

<u>AP</u>		BAXTER HLTHCARE	<u>10GM/100ML</u>	<u>N013684</u>	<u>006</u>	
-----------	--	-----------------	-------------------	----------------	------------	--

OSMITROL 15% IN WATER

<u>AP</u>		BAXTER HLTHCARE	<u>15GM/100ML</u>	<u>N013684</u>	<u>004</u>	
-----------	--	-----------------	-------------------	----------------	------------	--

OSMITROL 15% IN WATER IN PLASTIC CONTAINER

<u>AP</u>		BAXTER HLTHCARE	<u>15GM/100ML</u>	<u>N013684</u>	<u>008</u>	
-----------	--	-----------------	-------------------	----------------	------------	--

OSMITROL 20% IN WATER

<u>AP</u>		BAXTER HLTHCARE	<u>20GM/100ML</u>	<u>N013684</u>	<u>003</u>	
-----------	--	-----------------	-------------------	----------------	------------	--

OSMITROL 20% IN WATER IN PLASTIC CONTAINER

<u>AP</u>	+	!	BAXTER HLTHCARE	<u>20GM/100ML</u>	<u>N013684</u>	<u>007</u>
-----------	---	---	-----------------	-------------------	----------------	------------

OSMITROL 5% IN WATER

<u>AP</u>		BAXTER HLTHCARE	<u>5GM/100ML</u>	<u>N013684</u>	<u>001</u>	
-----------	--	-----------------	------------------	----------------	------------	--

OSMITROL 5% IN WATER IN PLASTIC CONTAINER

<u>AP</u>		BAXTER HLTHCARE	<u>5GM/100ML</u>	<u>N013684</u>	<u>005</u>	
-----------	--	-----------------	------------------	----------------	------------	--

POWDER; INHALATION

ARIDOL KIT

+	!	PHARMAXIS EUROPE	N/A, 5MG, 10MG, 20MG, 40MG	N022368	001	Oct 05, 2010
---	---	------------------	----------------------------	---------	-----	--------------

BRONCHITOL

+	!	PHARMAXIS EUROPE	40MG	N202049	001	Oct 30, 2020
---	---	------------------	------	---------	-----	--------------

MANNITOL; SORBITOL

SOLUTION; IRRIGATION

SORBITOL-MANNITOL IN PLASTIC CONTAINER

+	!	ICU MEDICAL INC	540MG/100ML; 2.7GM/100ML	N018316	001	
---	---	-----------------	--------------------------	---------	-----	--

MARALIXIBAT CHLORIDE

SOLUTION; ORAL

LIVMARLI

+	!	MIRUM	EQ 9.5MG BASE/ML	N214662	001	Sep 29, 2021
---	---	-------	------------------	---------	-----	--------------

+	!		EQ 19MG BASE/ML	N214662	002	Jul 24, 2024
---	---	--	-----------------	---------	-----	--------------

MARAVIROC

SOLUTION; ORAL

SELZENTRY

+	!	VIIV HLTHCARE	20MG/ML	N208984	001	Nov 04, 2016
---	---	---------------	---------	---------	-----	--------------

TABLET; ORAL

MARAVIROC

<u>AB</u>		HETERO LABS LTD III	<u>150MG</u>	<u>A203347</u>	<u>001</u>	Feb 07, 2022
-----------	--	---------------------	--------------	----------------	------------	--------------

<u>AB</u>			<u>300MG</u>	<u>A203347</u>	<u>002</u>	Feb 07, 2022
-----------	--	--	--------------	----------------	------------	--------------

<u>AB</u>		I 3 PHARMS	<u>150MG</u>	<u>A217114</u>	<u>001</u>	Aug 17, 2023
-----------	--	------------	--------------	----------------	------------	--------------

<u>AB</u>			<u>300MG</u>	<u>A217114</u>	<u>002</u>	Aug 17, 2023
-----------	--	--	--------------	----------------	------------	--------------

SELZENTRY

<u>AB</u>	+		<u>150MG</u>	<u>N022128</u>	<u>001</u>	Aug 06, 2007
-----------	---	--	--------------	----------------	------------	--------------

<u>AB</u>	+	!	<u>300MG</u>	<u>N022128</u>	<u>002</u>	Aug 06, 2007
-----------	---	---	--------------	----------------	------------	--------------

MARIBAVIR

TABLET; ORAL

LIVTENCITY

+	!	TAKEDA PHARMS USA	200MG	N215596	001	Nov 23, 2021
---	---	-------------------	-------	---------	-----	--------------

PRESCRIPTION DRUG PRODUCT LIST

MAVACAMTEN

CAPSULE; ORAL

CAMZYOS

+	BRISTOL	2.5MG	N214998	001	Apr 28, 2022
+	!	5MG	N214998	002	Apr 28, 2022
+		10MG	N214998	003	Apr 28, 2022
+		15MG	N214998	004	Apr 28, 2022

MAVORIXAFOR

CAPSULE; ORAL

XOLREMDI

+	X4 PHARMS	100MG	N218709	001	Apr 26, 2024
---	-----------	-------	---------	-----	--------------

MEBENDAZOLE

TABLET, CHEWABLE; ORAL

EMVERM

!	IMPAX LABS INC	100MG	A073580	001	Jan 04, 1995
---	----------------	-------	---------	-----	--------------

MECAMYLAMINE HYDROCHLORIDE

TABLET; ORAL

MECAMYLAMINE HYDROCHLORIDE

!	LGM PHARMA	2.5MG	A204054	001	Mar 19, 2013
---	------------	-------	---------	-----	--------------

MECHLORETHAMINE HYDROCHLORIDE

GEL; TOPICAL

VALCHLOR

+	HELSINN	EQ 0.016% BASE	N202317	001	Aug 23, 2013
---	---------	----------------	---------	-----	--------------

MECLIZINE HYDROCHLORIDE

TABLET; ORAL

ANTIVERT

<u>AA</u>	+	CASPER PHARMA LLC	<u>12.5MG</u>	<u>N010721</u>	<u>006</u>
<u>AA</u>	+		<u>25MG</u>	<u>N010721</u>	<u>004</u>
<u>AA</u>	+	!	<u>50MG</u>	<u>N010721</u>	<u>001</u> Jan 20, 1982

MECLIZINE HYDROCHLORIDE

<u>AA</u>	!	AMNEAL PHARMS	<u>12.5MG</u>	<u>A201451</u>	<u>001</u> Feb 23, 2011
<u>AA</u>	!		<u>25MG</u>	<u>A201451</u>	<u>002</u> Feb 23, 2011
<u>AA</u>			<u>50MG</u>	<u>A201451</u>	<u>003</u> Feb 23, 2011
<u>AA</u>		APNAR PHARMA LP	<u>12.5MG</u>	<u>A087128</u>	<u>002</u>
<u>AA</u>			<u>25MG</u>	<u>A087128</u>	<u>001</u>
<u>AA</u>		AUROBINDO PHARMA USA	<u>12.5MG</u>	<u>A202640</u>	<u>001</u> Sep 17, 2012
<u>AA</u>			<u>25MG</u>	<u>A202640</u>	<u>002</u> Sep 17, 2012
<u>AA</u>			<u>50MG</u>	<u>A202640</u>	<u>003</u> Sep 17, 2012
<u>AA</u>		CHARTWELL RX	<u>12.5MG</u>	<u>A203003</u>	<u>001</u> Aug 11, 2022
<u>AA</u>			<u>25MG</u>	<u>A203003</u>	<u>002</u> Aug 11, 2022
<u>AA</u>			<u>50MG</u>	<u>A203003</u>	<u>003</u> Aug 11, 2022
<u>AA</u>		EPIC PHARMA LLC	<u>12.5MG</u>	<u>A200294</u>	<u>001</u> Apr 13, 2012
<u>AA</u>			<u>25MG</u>	<u>A200294</u>	<u>002</u> Apr 13, 2012
<u>AA</u>		INVATECH	<u>25MG</u>	<u>A084092</u>	<u>003</u> May 22, 1989
<u>AA</u>		JUBILANT CADISTA	<u>12.5MG</u>	<u>A040659</u>	<u>001</u> Jun 04, 2010
<u>AA</u>			<u>25MG</u>	<u>A040659</u>	<u>002</u> Jun 04, 2010
<u>AA</u>		SANDOZ	<u>12.5MG</u>	<u>A084843</u>	<u>002</u> May 22, 1989
<u>AA</u>		ZYDUS LIFESCIENCES	<u>12.5MG</u>	<u>A213957</u>	<u>001</u> Jun 23, 2020
<u>AA</u>			<u>25MG</u>	<u>A213957</u>	<u>002</u> Jun 23, 2020

TABLET, CHEWABLE; ORAL

ANTIVERT

+	CASPER PHARMA LLC	25MG	N010721	005	
---	-------------------	------	---------	-----	--

MECLOFENAMATE SODIUM

CAPSULE; ORAL

MECLOFENAMATE SODIUM

	MYLAN	EQ 50MG BASE	A071081	002	Sep 03, 1986
!		EQ 100MG BASE	A071081	001	Sep 03, 1986

MEDROXYPROGESTERONE ACETATE

INJECTABLE; INJECTION

DEPO-PROVERA

<u>AB</u>	+	!	PFIZER	<u>150MG/ML</u>	<u>N020246</u>	<u>001</u> Oct 29, 1992
-----------	---	---	--------	-----------------	----------------	-------------------------

MEDROXYPROGESTERONE ACETATE

<u>AB</u>		AMNEAL	<u>150MG/ML</u>	<u>A215397</u>	<u>001</u> Jun 07, 2023
<u>AB</u>		AMPHASTAR PHARMS INC	<u>150MG/ML</u>	<u>A077235</u>	<u>001</u> Nov 28, 2017
<u>AB</u>			<u>150MG/ML</u>	<u>A077334</u>	<u>001</u> Nov 28, 2017
<u>AB</u>		EUGIA PHARMA	<u>150MG/ML</u>	<u>A212824</u>	<u>001</u> Aug 22, 2022
<u>AB</u>			<u>150MG/ML</u>	<u>A212844</u>	<u>001</u> Aug 31, 2022

PRESCRIPTION DRUG PRODUCT LIST

MEDROXYPROGESTERONE ACETATE

INJECTABLE; INJECTION

MEDROXYPROGESTERONE ACETATE

AB	HONG KONG	150MG/ML	A076553 001	Jul 28, 2004
AB	XIROMED	150MG/ML	A210227 001	Oct 12, 2018

INJECTABLE; SUBCUTANEOUS

DEPO-SUBQ PROVERA 104

+! PFIZER

104MG/0.65ML

N021583 001 Dec 17, 2004

TABLET; ORAL

MEDROXYPROGESTERONE ACETATE

AB	BARR	2.5MG	A040159 001	Aug 09, 1996
AB		5MG	A040159 002	Aug 09, 1996
AB		10MG	A040159 003	Aug 09, 1996

PROVERA

AB	+	PFIZER	2.5MG	N011839 001
AB	+		5MG	N011839 003
AB	+!		10MG	N011839 004

MEFENAMIC ACID

CAPSULE; ORAL

MEFENAMIC ACID

AB	BELCHER	250MG	A091608 001	Jun 02, 2014
AB	LUPIN LTD	250MG	A091322 001	Jul 22, 2011
AB	MICRO LABS	250MG	A090562 001	Nov 19, 2010

PONSTEL

AB	+!	AVION PHARMS	250MG	N015034 003
-----------	----	--------------	--------------	--------------------

MEFLOQUINE HYDROCHLORIDE

TABLET; ORAL

MEFLOQUINE HYDROCHLORIDE

AB	!	BARR	250MG	A076392 001	Dec 29, 2003
AB		HIKMA	250MG	A076523 001	Oct 01, 2004

MEGESTROL ACETATE

SUSPENSION; ORAL

MEGACE ES

AB	+!	ENDO OPERATIONS	125MG/ML	N021778 001	Jul 05, 2005
-----------	----	-----------------	-----------------	--------------------	--------------

MEGESTROL ACETATE

AB		NOVITIUM PHARMA	40MG/ML	A077404 001	Feb 16, 2006
AB	!	STRIDES PHARMA	40MG/ML	A075671 001	Jul 25, 2001
AB		TWI PHARMS	125MG/ML	A203139 001	Aug 27, 2014

TABLET; ORAL

MEGESTROL ACETATE

AB		BARR	20MG	A074621 002	Aug 16, 1996
AB			40MG	A074621 001	Nov 30, 1995
AB		STRIDES PHARMA	20MG	A072422 001	Aug 08, 1988
AB	!		40MG	A072423 001	Aug 08, 1988

MELOXICAM

CAPSULE; ORAL

MELOXICAM

AB		LUPIN LTD	5MG	A209487 001	Jun 01, 2020
AB	!		10MG	A209487 002	Jun 01, 2020
AB		NOVITIUM PHARMA	5MG	A211398 001	Mar 09, 2021
AB			10MG	A211398 002	Mar 09, 2021

SUSPENSION; ORAL

MELOXICAM

+! AVONDALE PHARMS

7.5MG/5ML

N021530 001 Jun 01, 2004

TABLET; ORAL

MELOXICAM

AB	AIPING PHARM INC	7.5MG	A077920 001	Jul 19, 2006
AB		15MG	A077920 002	Jul 19, 2006
AB	ASCENT PHARMS INC	7.5MG	A217579 001	Jun 07, 2023
AB		15MG	A217579 002	Jun 07, 2023
AB	AUROBINDO PHARMA	7.5MG	A078008 001	Oct 02, 2006
AB		15MG	A078008 002	Oct 02, 2006
AB	CIPLA	7.5MG	A077929 001	Jul 19, 2006
AB		15MG	A077929 002	Jul 19, 2006
AB	COREPHARMA	7.5MG	A077882 001	Jul 20, 2006
AB		15MG	A077882 002	Jul 20, 2006
AB	DR REDDYS LABS INC	7.5MG	A077931 001	Jul 25, 2006
AB		15MG	A077931 002	Jul 25, 2006
AB	ESJAY PHARMA	7.5MG	A077928 001	May 13, 2009
AB		15MG	A077928 002	May 13, 2009

PRESCRIPTION DRUG PRODUCT LIST

MELOXICAM

TABLET; ORAL

MELOXICAM

<u>AB</u>	GLENMARK PHARMS LTD	<u>7.5MG</u>	<u>A077932 001</u>	Jul 19, 2006
<u>AB</u>		<u>15MG</u>	<u>A077932 002</u>	Jul 19, 2006
<u>AB</u>	LUPIN PHARMS	<u>7.5MG</u>	<u>A077944 001</u>	Jul 19, 2006
<u>AB</u>		<u>15MG</u>	<u>A077944 002</u>	Jul 19, 2006
<u>AB</u>	PURACAP PHARM	<u>7.5MG</u>	<u>A077938 001</u>	Jul 19, 2006
<u>AB</u>		<u>15MG</u>	<u>A077938 002</u>	Jul 19, 2006
<u>AB</u>	TARO	<u>7.5MG</u>	<u>A078102 001</u>	Nov 07, 2006
<u>AB</u>		<u>15MG</u>	<u>A078102 002</u>	Nov 07, 2006
<u>AB</u>	UNICHEM	<u>7.5MG</u>	<u>A077927 001</u>	Dec 20, 2006
<u>AB</u>		<u>15MG</u>	<u>A077927 002</u>	Dec 20, 2006
<u>AB</u>	YUNG SHIN PHARM	<u>7.5MG</u>	<u>A077918 001</u>	Dec 07, 2006
<u>AB</u>		<u>15MG</u>	<u>A077918 002</u>	Dec 07, 2006
<u>AB</u>	ZYDUS PHARMS USA	<u>7.5MG</u>	<u>A077921 001</u>	Jul 19, 2006
<u>AB</u>	!	<u>15MG</u>	<u>A077921 002</u>	Jul 19, 2006

MELPHALAN HYDROCHLORIDE

INJECTABLE; INJECTION

MELPHALAN HYDROCHLORIDE

<u>AP</u>	ACTAVIS LLC	<u>EQ 50MG BASE/VIAL</u>	<u>A206018 001</u>	Dec 19, 2016
<u>AP</u>	ALMAJECT	<u>EQ 50MG BASE/VIAL</u>	<u>A204817 001</u>	May 17, 2019
<u>AP</u>	ARTHUR GRP	<u>EQ 50MG BASE/VIAL</u>	<u>A211463 001</u>	Sep 13, 2019
<u>AP</u>	AVET LIFESCIENCES	<u>EQ 50MG BASE/VIAL</u>	<u>A206523 001</u>	Oct 29, 2024
<u>AP</u>	BPI LABS	<u>EQ 50MG BASE/VIAL</u>	<u>A209197 001</u>	May 08, 2020
<u>AP</u>	DR REDDYS	<u>EQ 50MG BASE/VIAL</u>	<u>A203655 001</u>	Dec 08, 2017
<u>AP</u>	FRESENIUS KABI USA	<u>EQ 50MG BASE/VIAL</u>	<u>A203393 001</u>	Dec 22, 2017
<u>AP</u>	GLAND PHARMA LTD	<u>EQ 50MG BASE/VIAL</u>	<u>A209826 001</u>	May 28, 2019
<u>AP</u>	HETERO LABS	<u>EQ 50MG BASE/VIAL</u>	<u>A215024 001</u>	May 23, 2024
<u>AP</u>	HIKMA	<u>EQ 50MG BASE/VIAL</u>	<u>A090303 001</u>	Oct 28, 2010
<u>AP</u>	INGENUS PHARMS LLC	<u>EQ 50MG BASE/VIAL</u>	<u>A210947 001</u>	Feb 18, 2020
<u>AP</u>	MEITHEAL	<u>EQ 50MG BASE/VIAL</u>	<u>A212960 001</u>	May 28, 2021
<u>AP</u>	!	<u>EQ 50MG BASE/VIAL</u>	<u>A090270 001</u>	Jun 09, 2009
<u>AP</u>	SAGENT PHARMS INC	<u>EQ 50MG BASE/VIAL</u>	<u>A201379 001</u>	Feb 28, 2017

POWDER; INTRA-ARTERIAL

HEPZATO

+! DELCATH SYSTEMS INC EQ 50MG BASE/VIAL N201848 001 Aug 14, 2023

POWDER; INTRAVENOUS

EVOMELA

+! ACROTECH BIOPHARMA EQ 50MG BASE/VIAL N207155 001 Mar 10, 2016

SOLUTION; INTRAVENOUS

IVRA

+! APOTEX EQ 90MG BASE/ML (EQ 90MG BASE/ML) N217110 001 Aug 18, 2023

MEMANTINE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

MEMANTINE HYDROCHLORIDE

<u>AB</u>	AMNEAL PHARMS	<u>7MG</u>	<u>A205825 001</u>	Oct 12, 2016
<u>AB</u>		<u>14MG</u>	<u>A205825 002</u>	Oct 12, 2016
<u>AB</u>		<u>21MG</u>	<u>A205825 003</u>	Oct 12, 2016
<u>AB</u>		<u>28MG</u>	<u>A205825 004</u>	Oct 12, 2016
<u>AB</u>	APOTEX	<u>7MG</u>	<u>A206135 001</u>	Nov 22, 2016
<u>AB</u>		<u>14MG</u>	<u>A206135 002</u>	Nov 22, 2016
<u>AB</u>		<u>21MG</u>	<u>A206135 003</u>	Nov 22, 2016
<u>AB</u>		<u>28MG</u>	<u>A206135 004</u>	Nov 22, 2016
<u>AB</u>	AUROBINDO PHARMA LTD	<u>7MG</u>	<u>A214651 001</u>	Aug 09, 2021
<u>AB</u>		<u>14MG</u>	<u>A214651 002</u>	Aug 09, 2021
<u>AB</u>		<u>21MG</u>	<u>A214651 003</u>	Aug 09, 2021
<u>AB</u>		<u>28MG</u>	<u>A214651 004</u>	Aug 09, 2021
<u>AB</u>	LUPIN LTD	<u>7MG</u>	<u>A206028 001</u>	Sep 28, 2016
<u>AB</u>		<u>14MG</u>	<u>A206028 002</u>	Sep 28, 2016
<u>AB</u>		<u>21MG</u>	<u>A206028 003</u>	Sep 28, 2016
<u>AB</u>		<u>28MG</u>	<u>A206028 004</u>	Sep 28, 2016
<u>AB</u>	XIAMEN LP PHARM CO	<u>7MG</u>	<u>A213985 001</u>	Oct 11, 2022
<u>AB</u>		<u>14MG</u>	<u>A213985 002</u>	Oct 11, 2022
<u>AB</u>		<u>21MG</u>	<u>A213985 004</u>	Feb 06, 2024
<u>AB</u>		<u>28MG</u>	<u>A213985 003</u>	Oct 11, 2022
<u>AB</u>	YICHANG HUMANWELL	<u>7MG</u>	<u>A211100 001</u>	Apr 02, 2021
<u>AB</u>		<u>14MG</u>	<u>A211100 002</u>	Apr 02, 2021
<u>AB</u>		<u>21MG</u>	<u>A211100 003</u>	Apr 02, 2021
<u>AB</u>		<u>28MG</u>	<u>A211100 004</u>	Apr 02, 2021
<u>AB</u>	ZYDUS PHARMS	<u>7MG</u>	<u>A203293 001</u>	Aug 03, 2017

PRESCRIPTION DRUG PRODUCT LIST

MEMANTINE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE;ORAL

MEMANTINE HYDROCHLORIDE

<u>AB</u>		<u>14MG</u>	<u>A203293</u>	<u>002</u>	Aug 03, 2017
<u>AB</u>		<u>21MG</u>	<u>A203293</u>	<u>003</u>	Aug 03, 2017
<u>AB</u>	!	<u>28MG</u>	<u>A203293</u>	<u>004</u>	Aug 03, 2017

SOLUTION;ORAL

MEMANTINE HYDROCHLORIDE

<u>AA</u>	APOTEX	<u>2MG/ML</u>	<u>A209955</u>	<u>001</u>	Feb 09, 2018
<u>AA</u>	!	<u>2MG/ML</u>	<u>A204033</u>	<u>001</u>	Oct 13, 2015
<u>AA</u>	MACLEODS PHARMS LTD	<u>2MG/ML</u>	<u>A202790</u>	<u>001</u>	Oct 13, 2015
<u>AA</u>	SETON PHARMS	<u>2MG/ML</u>	<u>A210319</u>	<u>001</u>	Aug 31, 2020

TABLET;ORAL

MEMANTINE HYDROCHLORIDE

<u>AB</u>	AJANTA PHARMA LTD	<u>5MG</u>	<u>A206528</u>	<u>001</u>	Nov 30, 2015
<u>AB</u>		<u>10MG</u>	<u>A206528</u>	<u>002</u>	Nov 30, 2015
<u>AB</u>	ALEMBIC	<u>5MG</u>	<u>A200891</u>	<u>001</u>	Oct 13, 2015
<u>AB</u>		<u>10MG</u>	<u>A200891</u>	<u>002</u>	Oct 13, 2015
<u>AB</u>	AMNEAL PHARMS	<u>5MG</u>	<u>A090041</u>	<u>001</u>	Apr 10, 2015
<u>AB</u>		<u>10MG</u>	<u>A090041</u>	<u>002</u>	Apr 10, 2015
<u>AB</u>	AUROBINDO PHARMA LTD	<u>5MG</u>	<u>A203175</u>	<u>001</u>	Oct 13, 2015
<u>AB</u>		<u>10MG</u>	<u>A203175</u>	<u>002</u>	Oct 13, 2015
<u>AB</u>	DR REDDYS LABS LTD	<u>5MG</u>	<u>A090048</u>	<u>001</u>	Apr 14, 2010
<u>AB</u>		<u>10MG</u>	<u>A090048</u>	<u>002</u>	Apr 14, 2010
<u>AB</u>	HETERO LABS LTD V	<u>5MG</u>	<u>A204389</u>	<u>001</u>	Sep 26, 2022
<u>AB</u>		<u>10MG</u>	<u>A204389</u>	<u>002</u>	Sep 26, 2022
<u>AB</u>	LUPIN LTD	<u>5MG</u>	<u>A090051</u>	<u>001</u>	Apr 10, 2015
<u>AB</u>		<u>10MG</u>	<u>A090051</u>	<u>002</u>	Apr 10, 2015
<u>AB</u>	MACLEODS PHARMS LTD	<u>5MG</u>	<u>A202840</u>	<u>001</u>	Oct 13, 2015
<u>AB</u>		<u>10MG</u>	<u>A202840</u>	<u>002</u>	Oct 13, 2015
<u>AB</u>	POLYGEN PHARMS	<u>5MG</u>	<u>A210587</u>	<u>001</u>	Dec 11, 2020
<u>AB</u>		<u>10MG</u>	<u>A210587</u>	<u>002</u>	Dec 11, 2020
<u>AB</u>	PURACAP PHARM LLC	<u>5MG</u>	<u>A206855</u>	<u>001</u>	Nov 17, 2015
<u>AB</u>		<u>10MG</u>	<u>A206855</u>	<u>002</u>	Nov 17, 2015
<u>AB</u>	RENATA	<u>5MG</u>	<u>A209527</u>	<u>001</u>	May 07, 2018
<u>AB</u>		<u>10MG</u>	<u>A209527</u>	<u>002</u>	May 07, 2018
<u>AB</u>	STRIDES PHARMA	<u>5MG</u>	<u>A202350</u>	<u>001</u>	May 23, 2017
<u>AB</u>		<u>10MG</u>	<u>A202350</u>	<u>002</u>	May 23, 2017
<u>AB</u>	SUN PHARM	<u>5MG</u>	<u>A090058</u>	<u>001</u>	May 05, 2010
<u>AB</u>		<u>10MG</u>	<u>A090058</u>	<u>002</u>	May 05, 2010
<u>AB</u>	UNICHEM	<u>5MG</u>	<u>A200022</u>	<u>001</u>	Oct 13, 2015
<u>AB</u>		<u>10MG</u>	<u>A200022</u>	<u>002</u>	Oct 13, 2015
<u>AB</u>	UPSHER SMITH LABS	<u>5MG</u>	<u>A090043</u>	<u>001</u>	Jul 31, 2015
<u>AB</u>		<u>10MG</u>	<u>A090043</u>	<u>002</u>	Jul 31, 2015
<u>AB</u>	ZYDUS LIFESCIENCES	<u>5MG</u>	<u>A090961</u>	<u>001</u>	Jul 10, 2017
<u>AB</u>		<u>10MG</u>	<u>A090961</u>	<u>002</u>	Jul 10, 2017

NAMENDA

<u>AB</u>	+	ABEVIE	<u>5MG</u>	<u>N021487</u>	<u>001</u>	Oct 16, 2003
<u>AB</u>	+	!	<u>10MG</u>	<u>N021487</u>	<u>002</u>	Oct 16, 2003

MEPERIDINE HYDROCHLORIDE

INJECTABLE;INJECTION

DEMEROL

<u>AP</u>	+	!	HOSPIRA	<u>25MG/ML</u>	<u>N021171</u>	<u>001</u>
<u>AP</u>	+	!		<u>50MG/ML</u>	<u>N021171</u>	<u>002</u>
<u>AP</u>	+	!		<u>75MG/ML</u>	<u>N021171</u>	<u>003</u>
<u>AP</u>	+	!		<u>100MG/ML</u>	<u>N021171</u>	<u>004</u>

MEPERIDINE HYDROCHLORIDE

<u>AP</u>		WEST-WARD PHARMS INT	<u>25MG/ML</u>	<u>A080445</u>	<u>001</u>	
<u>AP</u>			<u>25MG/ML</u>	<u>A080455</u>	<u>007</u>	
<u>AP</u>			<u>50MG/ML</u>	<u>A080445</u>	<u>002</u>	
<u>AP</u>			<u>50MG/ML</u>	<u>A080455</u>	<u>008</u>	
<u>AP</u>			<u>75MG/ML</u>	<u>A080445</u>	<u>003</u>	
<u>AP</u>			<u>75MG/ML</u>	<u>A080455</u>	<u>009</u>	
<u>AP</u>			<u>100MG/ML</u>	<u>A080445</u>	<u>004</u>	
<u>AP</u>			<u>100MG/ML</u>	<u>A080455</u>	<u>010</u>	

MEPERIDINE HYDROCHLORIDE PRESERVATIVE FREE

!		WEST-WARD PHARMS INT	<u>10MG/ML</u>	<u>A081002</u>	<u>001</u>	Jul 30, 1993
---	--	----------------------	----------------	----------------	------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

MEPERIDINE HYDROCHLORIDE

SYRUP; ORAL

MEPERIDINE HYDROCHLORIDE

! HIKMA

50MG/5ML

A088744 001 Jan 30, 1985

TABLET; ORAL

MEPERIDINE HYDROCHLORIDEAA ! EPIC PHARMA LLC50MGA040331 001 May 28, 1999AA GENUS50MGA040893 001 Jun 24, 2009

EPIC PHARMA LLC

100MG

A040331 002 May 28, 1999

MEPIVACAINE HYDROCHLORIDE

INJECTABLE; INJECTION

CARBOCAINEAP +! HOSPIRA1%N012250 001AP +!1.5%N012250 005AP +!2%N012250 002POLOCAINEAP FRESENIUS KABI USA1%A089407 001 Dec 01, 1986AP2%A089410 001 Dec 01, 1986POLOCAINE-MPFAP FRESENIUS KABI USA1%A089406 001 Dec 01, 1986AP1.5%A089408 001 Dec 01, 1986AP2%A089409 001 Dec 01, 1986

SCANDONEST PLAIN

! DEPROCO

3%

A088387 001 Oct 10, 1984

MEPROBAMATE

TABLET; ORAL

MEPROBAMATEAA ! ALEMBIC PHARMS LTD200MGA090122 001 Feb 18, 2009AA !400MGA090122 002 Feb 18, 2009AA INVAGEN PHARMS200MGA040797 001 Feb 27, 2008AA400MGA040797 002 Feb 27, 2008MERCAPTOPYRINE

SUSPENSION; ORAL

PURIXAN

+! NOVA LABS LTD

20MG/ML

N205919 001 Apr 28, 2014

TABLET; ORAL

MERCAPTOPYRINEAB DR REDDYS LABS SA50MGA040461 001 Feb 11, 2004AB !

HIKMA

50MGA040528 001 Feb 13, 2004AB MYLAN50MGA040594 001 Jul 01, 2005PURINETHOLAB + STASON PHARMS50MGN009053 002MEROPEM

INJECTABLE; INJECTION

MEROPEMAP ACS DOBFAR500MG/VIALA091404 001 Oct 26, 2011AP1GM/VIALA091404 002 Oct 26, 2011AP ACS DOBFAR SPA500MG/VIALA204139 001 Jun 09, 2016AP1GM/VIALA204139 002 Jun 09, 2016AP AMNEAL PHARMS500MG/VIALA205883 001 Apr 12, 2016AP1GM/VIALA205883 002 Apr 12, 2016AP BROOKS STERISCIENCE500MG/VIALA216154 001 Aug 18, 2022AP1GM/VIALA216154 002 Aug 18, 2022AP EUGIA PHARMA500MG/VIALA205835 001 Mar 27, 2017AP1GM/VIALA205835 002 Mar 27, 2017AP GLAND500MG/VIALA206141 001 Jun 08, 2016AP1GM/VIALA206141 002 Jun 08, 2016AP HQ SPCLT PHARMA500MG/VIALA210773 001 Aug 16, 2019AP1GM/VIALA210773 002 Aug 16, 2019AP QILU500MG/VIALA216424 001 Jul 22, 2024AP1GM/VIALA216424 002 Jul 22, 2024AP SAVIOR LIFETEC CORP500MG/VIALA206086 001 Apr 19, 2016AP1GM/VIALA206086 002 Apr 19, 2016MERREM IVAP +! PFIZER500MG/VIALN050706 003 Jun 21, 1996AP +!1GM/VIALN050706 001 Jun 21, 1996

POWDER; INTRAVENOUS

MEROPEM

+! HQ SPCLT PHARMA

2GM/VIAL

N215212 001 Jul 26, 2023

PRESCRIPTION DRUG PRODUCT LIST

MEROPENEM

POWDER; INTRAVENOUS

MEROPENEM AND SODIUM CHLORIDE IN DUPLEX CONTAINER

B BRAUN MEDICAL INC 500MG/VIAL
1GM/VIALN202106 001 Apr 30, 2015
N202106 002 Apr 30, 2015MEROPENEM; VABORBACTAM

POWDER; INTRAVENOUS

VABOMERE

+! REMPEX 1GM/VIAL; 1GM/VIAL

N209776 001 Aug 29, 2017

MESALAMINE

CAPSULE, DELAYED RELEASE; ORAL

MESALAMINE

! TEVA PHARMS USA 400MG

A207873 001 May 09, 2019

CAPSULE, EXTENDED RELEASE; ORAL

APRISOAB +! SALIX375MGN022301 001 Oct 31, 2008MESALAMINEAB ALEMBIC375MGA216967 001 Oct 28, 2022AB ALKEM LABS LTD375MGA214242 001 Jul 15, 2021AB AMTA375MGA217533 001 Jun 06, 2023AB AUROBINDO PHARMA
LTD375MGA214477 001 Mar 31, 2023AB MYLAN375MGA207271 001 Nov 20, 2019AB NOVAST LABS375MGA218410 001 Aug 22, 2024AB SUN PHARM500MGA214585 001 May 11, 2022AB ZYDUS PHARMS375MGA208954 001 Aug 12, 2021PENTASAAB +! TAKEDA PHARMS USA500MGN020049 002 Jul 08, 2004

+

250MG

N020049 001 May 10, 1993

ENEMA; RECTAL

MESALAMINEAB ENCUBE4GM/60MLA216941 001 May 30, 2023AB PADAGIS ISRAEL4GM/60MLA076751 001 Sep 17, 2004ROWASAAB +! MYLAN SPECIALITY LP4GM/60MLN019618 001 Dec 24, 1987SFROWASAAB + MYLAN SPECIALITY LP4GM/60MLN019618 002 Jun 20, 2008

SUPPOSITORY; RECTAL

CANASAAB +! ABBEVIE1GMN021252 002 Nov 05, 2004MESALAMINEAB ACTAVIS MID1GMA205654 001 Aug 14, 2020

ATLANTIC

AB AMRING PHARMS1GMA208362 001 Jun 21, 2019AB ANNORA PHARMA1GMA213377 001 Mar 19, 2020AB MYLAN1GMA204354 001 Nov 24, 2015AB RISING1GMA207448 001 Apr 19, 2019AB SANDOZ1GMA202065 001 Jun 12, 2019AB ZYDUS PHARMS1GMA208953 001 Feb 12, 2020

TABLET, DELAYED RELEASE; ORAL

LIALDAAB +! TAKEDA PHARMS USA1.2GMN022000 001 Jan 16, 2007MESALAMINEAB ACTAVIS LABS FL1.2GMA203817 001 Mar 23, 2018AB ANNORA PHARMA1.2GMA216334 001 Feb 05, 2024AB SINOTHERAPEUTICS1.2GMA217337 001 May 12, 2023

INC

AB SUN PHARM1.2GMA211858 001 Jan 25, 2019AB TEVA PHARMS INC800MGA213191 001 Aug 22, 2024AB ! ZYDUS PHARMS800MGA203286 001 Jul 21, 2017AB 1.2GM A091640 001 Jun 05, 2017MESNA

INJECTABLE; INTRAVENOUS

MESNAAP FRESENIUS KABI USA100MG/MLA075811 001 Apr 26, 2001AP GLAND100MG/MLA206992 001 Dec 18, 2017AP HIKMA100MG/MLA075739 001 Jan 09, 2004AP SAGENT PHARMS INC100MG/MLA090913 001 Apr 13, 2010MESNEXAP +! BAXTER HLTHCARE100MG/MLN019884 001 Dec 30, 1988

PRESCRIPTION DRUG PRODUCT LIST

MESNA

TABLET; ORAL

MESNEX

+! BAXTER HLTHCARE 400MG N020855 001 Mar 21, 2002

METAPROTERENOL SULFATE

SYRUP; ORAL

METAPROTERENOL SULFATE

! GENUS LIFESCIENCES 10MG/5ML A073632 001 Jul 22, 1992

METARAMINOL BITARTRATE

INJECTABLE; INJECTION

METARAMINOL BITARTRATE

AZURITY EQ 10MG BASE/ML A211304 001 Aug 24, 2021

METAXALONE

TABLET; ORAL

METAXALONE

<u>AB</u>	ACTAVIS LABS FL INC	<u>800MG</u>	<u>A203695</u>	<u>001</u>	Jun 15, 2017
<u>AB</u>	! AMNEAL PHARMS	<u>800MG</u>	<u>A203399</u>	<u>001</u>	Jun 21, 2013
<u>AB</u>	LANNETT CO INC	<u>800MG</u>	<u>A204770</u>	<u>001</u>	Nov 22, 2016
<u>AB</u>	MOUNTAIN	<u>400MG</u>	<u>A040486</u>	<u>001</u>	Feb 27, 2015
<u>AB</u>	SANDOZ	<u>800MG</u>	<u>A040445</u>	<u>001</u>	Mar 31, 2010
<u>AB</u>	SCIEGEN PHARMS INC	<u>400MG</u>	<u>A207466</u>	<u>002</u>	Mar 13, 2020
<u>AB</u>		<u>800MG</u>	<u>A207466</u>	<u>001</u>	Aug 31, 2017

METFORMIN HYDROCHLORIDE

SOLUTION; ORAL

METFORMIN HYDROCHLORIDE

<u>AB</u>	! SAPTALIS PHARMS	<u>500MG/5ML</u>	<u>A211309</u>	<u>001</u>	Mar 03, 2020
<u>AB</u>	VISTAPHARM	<u>500MG/5ML</u>	<u>A212677</u>	<u>001</u>	Aug 19, 2022

TABLET; ORAL

METFORMIN HYDROCHLORIDE

<u>AB</u>	ALKEM	<u>500MG</u>	<u>A091184</u>	<u>001</u>	Nov 01, 2010
<u>AB</u>		<u>850MG</u>	<u>A091184</u>	<u>002</u>	Nov 01, 2010
<u>AB</u>		<u>1GM</u>	<u>A091184</u>	<u>003</u>	Nov 01, 2010
<u>AB</u>	APOTEX	<u>500MG</u>	<u>A075984</u>	<u>001</u>	Apr 23, 2002
<u>AB</u>		<u>500MG</u>	<u>A090666</u>	<u>001</u>	Dec 07, 2011
<u>AB</u>		<u>850MG</u>	<u>A075984</u>	<u>002</u>	Apr 23, 2002
<u>AB</u>		<u>850MG</u>	<u>A090666</u>	<u>002</u>	Dec 07, 2011
<u>AB</u>		<u>1GM</u>	<u>A075984</u>	<u>003</u>	Apr 23, 2002
<u>AB</u>		<u>1GM</u>	<u>A090666</u>	<u>003</u>	Dec 07, 2011
<u>AB</u>	ATLAS PHARMS LLC	<u>500MG</u>	<u>A076033</u>	<u>001</u>	Jan 24, 2002
<u>AB</u>		<u>850MG</u>	<u>A076033</u>	<u>002</u>	Jan 24, 2002
<u>AB</u>		<u>1GM</u>	<u>A076033</u>	<u>003</u>	Jan 24, 2002
<u>AB</u>	AUROBINDO	<u>500MG</u>	<u>A077095</u>	<u>001</u>	Jan 14, 2005
<u>AB</u>		<u>850MG</u>	<u>A077095</u>	<u>002</u>	Jan 14, 2005
<u>AB</u>		<u>1GM</u>	<u>A077095</u>	<u>003</u>	Jan 14, 2005
<u>AB</u>	CHARTWELL	<u>500MG</u>	<u>A075972</u>	<u>001</u>	Jan 24, 2002
<u>AB</u>		<u>850MG</u>	<u>A075972</u>	<u>002</u>	Jan 24, 2002
<u>AB</u>		<u>1GM</u>	<u>A075972</u>	<u>003</u>	Jan 24, 2002
<u>AB</u>	CSPC OUYI	<u>500MG</u>	<u>A205096</u>	<u>001</u>	Jul 11, 2016
<u>AB</u>		<u>850MG</u>	<u>A205096</u>	<u>002</u>	Jul 11, 2016
<u>AB</u>		<u>1GM</u>	<u>A205096</u>	<u>003</u>	Jul 11, 2016
<u>AB</u>	DR REDDYS LABS INC	<u>500MG</u>	<u>A077787</u>	<u>001</u>	Aug 23, 2006
<u>AB</u>		<u>850MG</u>	<u>A077787</u>	<u>002</u>	Aug 23, 2006
<u>AB</u>		<u>1GM</u>	<u>A077787</u>	<u>003</u>	Aug 23, 2006
<u>AB</u>	EPIC PHARMA LLC	<u>500MG</u>	<u>A075965</u>	<u>001</u>	Jan 25, 2002
<u>AB</u>		<u>850MG</u>	<u>A075965</u>	<u>002</u>	Jan 25, 2002
<u>AB</u>		<u>1GM</u>	<u>A075965</u>	<u>003</u>	Jan 25, 2002
<u>AB</u>	GLENMARK PHARMS LTD	<u>500MG</u>	<u>A078170</u>	<u>001</u>	May 23, 2008
<u>AB</u>		<u>850MG</u>	<u>A078170</u>	<u>002</u>	May 23, 2008
<u>AB</u>		<u>1GM</u>	<u>A078170</u>	<u>003</u>	May 23, 2008
<u>AB</u>	GRANULES INDIA	<u>500MG</u>	<u>A090564</u>	<u>001</u>	Apr 22, 2010
<u>AB</u>		<u>850MG</u>	<u>A090564</u>	<u>002</u>	Apr 22, 2010
<u>AB</u>	!	<u>1GM</u>	<u>A090564</u>	<u>003</u>	Apr 22, 2010
<u>AB</u>	HARMAN FINOCHEM	<u>500MG</u>	<u>A213320</u>	<u>001</u>	Dec 03, 2021
<u>AB</u>		<u>850MG</u>	<u>A213320</u>	<u>002</u>	Dec 03, 2021
<u>AB</u>		<u>1GM</u>	<u>A213320</u>	<u>003</u>	Dec 03, 2021
<u>AB</u>	LAURUS	<u>500MG</u>	<u>A209882</u>	<u>001</u>	Aug 27, 2018
<u>AB</u>		<u>850MG</u>	<u>A209882</u>	<u>002</u>	Aug 27, 2018
<u>AB</u>		<u>1GM</u>	<u>A209882</u>	<u>003</u>	Aug 27, 2018
<u>AB</u>	MARKSANS PHARMA	<u>500MG</u>	<u>A090888</u>	<u>001</u>	Mar 12, 2012
<u>AB</u>		<u>850MG</u>	<u>A090888</u>	<u>002</u>	Mar 12, 2012

PRESCRIPTION DRUG PRODUCT LIST

METFORMIN HYDROCHLORIDE

TABLET; ORAL

METFORMIN HYDROCHLORIDE

<u>AB</u>		<u>1GM</u>	<u>A090888</u>	<u>003</u>	Mar 12, 2012
<u>AB</u>	MYLAN	<u>500MG</u>	<u>A075973</u>	<u>001</u>	Jan 25, 2002
<u>AB</u>		<u>850MG</u>	<u>A075973</u>	<u>002</u>	Jan 25, 2002
<u>AB</u>		<u>1GM</u>	<u>A075973</u>	<u>003</u>	Jan 25, 2002
<u>AB</u>	SCIEGEN PHARMS INC	<u>500MG</u>	<u>A203769</u>	<u>001</u>	Sep 11, 2013
<u>AB</u>		<u>850MG</u>	<u>A203769</u>	<u>002</u>	Sep 11, 2013
<u>AB</u>		<u>1GM</u>	<u>A203769</u>	<u>003</u>	Sep 11, 2013
<u>AB</u>	ZYDUS HLTHCARE	<u>500MG</u>	<u>A203686</u>	<u>001</u>	Aug 28, 2014
<u>AB</u>		<u>850MG</u>	<u>A203686</u>	<u>002</u>	Aug 28, 2014
<u>AB</u>		<u>1GM</u>	<u>A203686</u>	<u>003</u>	Aug 28, 2014
<u>AB</u>	ZYDUS PHARMS USA	<u>500MG</u>	<u>A077064</u>	<u>001</u>	Apr 18, 2005
<u>AB</u>		<u>850MG</u>	<u>A077064</u>	<u>002</u>	Apr 18, 2005
<u>AB</u>		<u>1GM</u>	<u>A077064</u>	<u>003</u>	Apr 18, 2005
	CHARTWELL	625MG	A075972	005	Jan 24, 2002
		750MG	A075972	004	Jan 24, 2002

TABLET, EXTENDED RELEASE; ORAL

METFORMIN HYDROCHLORIDE

<u>AB</u>	ACTAVIS LABS FL INC	<u>750MG</u>	<u>A076869</u>	<u>001</u>	Apr 12, 2005
<u>AB</u>	! ALKEM LABS LTD	<u>750MG</u>	<u>A206145</u>	<u>002</u>	Oct 22, 2018
<u>AB</u>	AMNEAL PHARMS NY	<u>750MG</u>	<u>A078596</u>	<u>002</u>	Jan 03, 2008
<u>AB</u>	AUROBINDO PHARMA LTD	<u>750MG</u>	<u>A079118</u>	<u>002</u>	Jul 20, 2012
<u>AB</u>	BEXIMCO PHARMS USA	<u>750MG</u>	<u>A207427</u>	<u>002</u>	Dec 13, 2016
<u>AB</u>	CSPC OUYI	<u>750MG</u>	<u>A078321</u>	<u>002</u>	Apr 17, 2008
<u>AB</u>	GRANULES	<u>750MG</u>	<u>A209313</u>	<u>002</u>	Mar 16, 2018
<u>AB</u>	HARMAN FINOCHEM	<u>750MG</u>	<u>A218673</u>	<u>002</u>	Jul 05, 2024
<u>AB</u>	INTELLIPHARMACEUTICS	<u>750MG</u>	<u>A202306</u>	<u>002</u>	Feb 23, 2017
<u>AB</u>	LAURUS	<u>750MG</u>	<u>A217631</u>	<u>002</u>	Oct 05, 2023
<u>AB</u>	MACLEODS PHARMS LTD	<u>750MG</u>	<u>A206955</u>	<u>002</u>	Dec 07, 2016
<u>AB</u>	MARKSANS PHARMA	<u>750MG</u>	<u>A090295</u>	<u>002</u>	Apr 29, 2016
<u>AB</u>	NOSTRUM PHARMS LLC	<u>750MG</u>	<u>A076756</u>	<u>002</u>	Dec 12, 2011
<u>AB</u>	PRINSTON INC	<u>750MG</u>	<u>A208880</u>	<u>002</u>	Sep 10, 2018
<u>AB</u>	SUN PHARM INDS (IN)	<u>750MG</u>	<u>A077336</u>	<u>002</u>	Feb 09, 2006
<u>AB</u>	TEVA	<u>750MG</u>	<u>A076864</u>	<u>001</u>	Apr 12, 2005
<u>AB</u>	UNICHEM	<u>750MG</u>	<u>A213359</u>	<u>002</u>	Aug 11, 2021
<u>AB</u>	YICHANG HUMANWELL	<u>750MG</u>	<u>A211052</u>	<u>002</u>	Sep 24, 2018
<u>AB</u>	ZYDUS LIFESCIENCES	<u>750MG</u>	<u>A077078</u>	<u>001</u>	Apr 21, 2005
<u>AB1</u>	ALIGNSCIENCE PHARMA	<u>500MG</u>	<u>A209303</u>	<u>001</u>	Mar 19, 2018
<u>AB1</u>	ALKEM LABS LTD	<u>500MG</u>	<u>A206145</u>	<u>001</u>	Oct 22, 2018
<u>AB1</u>	AMNEAL PHARMS NY	<u>500MG</u>	<u>A078596</u>	<u>001</u>	Jan 03, 2008
<u>AB1</u>	AUROBINDO PHARMA LTD	<u>500MG</u>	<u>A079118</u>	<u>001</u>	Jul 20, 2012
<u>AB1</u>	BEXIMCO PHARMS USA	<u>500MG</u>	<u>A207427</u>	<u>001</u>	Dec 13, 2016
<u>AB1</u>	CSPC OUYI	<u>500MG</u>	<u>A078321</u>	<u>001</u>	Apr 17, 2008
<u>AB1</u>	GRANULES	<u>500MG</u>	<u>A209313</u>	<u>001</u>	Mar 16, 2018
<u>AB1</u>	HARMAN FINOCHEM	<u>500MG</u>	<u>A218673</u>	<u>001</u>	Jul 05, 2024
<u>AB1</u>	INTELLIPHARMACEUTICS	<u>500MG</u>	<u>A202306</u>	<u>001</u>	Feb 23, 2017
<u>AB1</u>	INVENTIA	<u>500MG</u>	<u>A201991</u>	<u>001</u>	Jan 18, 2012
<u>AB1</u>	LAURUS	<u>500MG</u>	<u>A217631</u>	<u>001</u>	Oct 05, 2023
<u>AB1</u>	MACLEODS PHARMS LTD	<u>500MG</u>	<u>A206955</u>	<u>001</u>	Dec 07, 2016
<u>AB1</u>	MARKSANS PHARMA	<u>500MG</u>	<u>A090295</u>	<u>001</u>	Apr 29, 2016
<u>AB1</u>	NOSTRUM PHARMS LLC	<u>500MG</u>	<u>A076756</u>	<u>001</u>	Jul 26, 2006
<u>AB1</u>	PRINSTON INC	<u>500MG</u>	<u>A208880</u>	<u>001</u>	Sep 10, 2018
<u>AB1</u>	SANDOZ	<u>500MG</u>	<u>A076873</u>	<u>001</u>	Dec 14, 2004
<u>AB1</u>	SUN PHARM INDS (IN)	<u>500MG</u>	<u>A077336</u>	<u>001</u>	Feb 09, 2006
<u>AB1</u>	TEVA	<u>500MG</u>	<u>A076269</u>	<u>001</u>	Jun 18, 2004
<u>AB1</u>	UNICHEM	<u>500MG</u>	<u>A213359</u>	<u>001</u>	Aug 11, 2021
<u>AB1</u>	YICHANG HUMANWELL	<u>500MG</u>	<u>A211052</u>	<u>001</u>	Sep 24, 2018
<u>AB1</u>	ZYDUS LIFESCIENCES	<u>500MG</u>	<u>A077060</u>	<u>001</u>	Apr 20, 2005

FORTAMET

<u>AB2</u>	+ ANDRX LABS LLC	<u>1GM</u>	<u>N021574</u>	<u>002</u>	Apr 27, 2004
------------	------------------	------------	----------------	------------	--------------

METFORMIN HYDROCHLORIDE

<u>AB2</u>	AJANTA PHARMA LTD	<u>500MG</u>	<u>A213651</u>	<u>001</u>	Apr 09, 2020
<u>AB2</u>		<u>1GM</u>	<u>A213651</u>	<u>002</u>	Apr 09, 2020
<u>AB2</u>	AUROBINDO PHARMA	<u>500MG</u>	<u>A209694</u>	<u>001</u>	Oct 18, 2024
<u>AB2</u>		<u>1GM</u>	<u>A209694</u>	<u>002</u>	Oct 18, 2024
<u>AB2</u>	LUPIN LTD	<u>500MG</u>	<u>A090692</u>	<u>001</u>	Jun 29, 2011
<u>AB2</u>	!	<u>1GM</u>	<u>A090692</u>	<u>002</u>	Jun 29, 2011

PRESCRIPTION DRUG PRODUCT LIST

METFORMIN HYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL

METFORMIN HYDROCHLORIDE

<u>AB2</u>	MYLAN PHARMS INC	<u>500MG</u>	<u>A200690</u>	<u>001</u>	Aug 01, 2012
<u>AB2</u>		<u>1GM</u>	<u>A200690</u>	<u>002</u>	Aug 01, 2012
<u>AB2</u>	NOSTRUM LABS INC	<u>500MG</u>	<u>A203832</u>	<u>001</u>	Dec 26, 2017
<u>AB2</u>		<u>1GM</u>	<u>A203832</u>	<u>002</u>	Dec 26, 2017
<u>AB2</u>	NOVAST LABS	<u>500MG</u>	<u>A209674</u>	<u>001</u>	Nov 02, 2018
<u>AB2</u>		<u>1GM</u>	<u>A209674</u>	<u>002</u>	Nov 02, 2018
<u>AB2</u>	QINGDAO BAHEAL PHARM	<u>500MG</u>	<u>A209993</u>	<u>001</u>	Dec 27, 2018
<u>AB2</u>		<u>1GM</u>	<u>A209993</u>	<u>002</u>	Dec 27, 2018
<u>AB2</u>	SCIEGEN PHARMS INC	<u>500MG</u>	<u>A214629</u>	<u>001</u>	Feb 22, 2022
<u>AB2</u>		<u>1GM</u>	<u>A214629</u>	<u>002</u>	Feb 22, 2022
<u>AB2</u>	TWI PHARMS	<u>500MG</u>	<u>A213247</u>	<u>001</u>	Sep 29, 2021
<u>AB2</u>		<u>1GM</u>	<u>A213247</u>	<u>002</u>	Sep 29, 2021

GLUMETZA

<u>AB3</u>	+ SANTARUS INC	<u>500MG</u>	<u>N021748</u>	<u>001</u>	Jun 03, 2005
<u>AB3</u>	+!	<u>1GM</u>	<u>N021748</u>	<u>002</u>	Jun 03, 2005

METFORMIN HYDROCHLORIDE

<u>AB3</u>	ACTAVIS LABS FL INC	<u>500MG</u>	<u>A203755</u>	<u>001</u>	Aug 01, 2016
<u>AB3</u>		<u>1GM</u>	<u>A203755</u>	<u>002</u>	Aug 01, 2016
<u>AB3</u>	AJANTA PHARMA LTD	<u>500MG</u>	<u>A213962</u>	<u>001</u>	Mar 09, 2021
<u>AB3</u>		<u>1GM</u>	<u>A213962</u>	<u>002</u>	Mar 09, 2021
<u>AB3</u>	APOTEX	<u>500MG</u>	<u>A213356</u>	<u>001</u>	Dec 13, 2021
<u>AB3</u>		<u>1GM</u>	<u>A213356</u>	<u>002</u>	Dec 13, 2021
<u>AB3</u>	GLENMARK PHARMS LTD	<u>500MG</u>	<u>A212969</u>	<u>001</u>	Nov 25, 2019
<u>AB3</u>		<u>1GM</u>	<u>A212969</u>	<u>002</u>	Nov 25, 2019
<u>AB3</u>	GRANULES	<u>500MG</u>	<u>A213344</u>	<u>001</u>	Jan 12, 2021
<u>AB3</u>		<u>1GM</u>	<u>A213344</u>	<u>002</u>	Jan 12, 2021
<u>AB3</u>	LUPIN LTD	<u>500MG</u>	<u>A091664</u>	<u>001</u>	Jul 19, 2013
<u>AB3</u>		<u>1GM</u>	<u>A091664</u>	<u>002</u>	Jul 19, 2013
<u>AB3</u>	MICRO LABS	<u>500MG</u>	<u>A212448</u>	<u>001</u>	Jan 10, 2023
<u>AB3</u>		<u>1GM</u>	<u>A212448</u>	<u>002</u>	Jan 10, 2023
<u>AB3</u>	PRINSTON INC	<u>500MG</u>	<u>A212681</u>	<u>001</u>	Jun 09, 2022
<u>AB3</u>		<u>1GM</u>	<u>A212681</u>	<u>002</u>	Jun 09, 2022
<u>AB3</u>	RK PHARMA	<u>500MG</u>	<u>A215629</u>	<u>001</u>	Jun 20, 2023
<u>AB3</u>		<u>1GM</u>	<u>A215629</u>	<u>002</u>	Jun 20, 2023
<u>AB3</u>	SCIEGEN PHARMS INC	<u>500MG</u>	<u>A213334</u>	<u>001</u>	Apr 16, 2021
<u>AB3</u>		<u>1GM</u>	<u>A213334</u>	<u>002</u>	Apr 16, 2021
<u>AB3</u>	SUN PHARM	<u>1GM</u>	<u>A202917</u>	<u>002</u>	Aug 01, 2016
<u>AB3</u>		<u>500MG</u>	<u>A202917</u>	<u>001</u>	Aug 01, 2016

METFORMIN HYDROCHLORIDE; PIOGLITAZONE HYDROCHLORIDE

TABLET;ORAL

ACTOPLUS MET

<u>AB</u>	+! TAKEDA PHARMS USA	<u>850MG;EQ 15MG BASE</u>	<u>N021842</u>	<u>002</u>	Aug 29, 2005
-----------	----------------------	---------------------------	----------------	------------	--------------

PIOGLITAZONE HYDROCHLORIDE AND METFORMIN HYDROCHLORIDE

<u>AB</u>	AUROBINDO PHARMA LTD	<u>500MG;EQ 15MG BASE</u>	<u>A200823</u>	<u>001</u>	Feb 13, 2013
<u>AB</u>		<u>850MG;EQ 15MG BASE</u>	<u>A200823</u>	<u>002</u>	Feb 13, 2013
<u>AB</u>	MACLEODS PHARMS LTD	<u>500MG;EQ 15MG BASE</u>	<u>A204802</u>	<u>001</u>	Nov 05, 2015
<u>AB</u>		<u>850MG;EQ 15MG BASE</u>	<u>A204802</u>	<u>002</u>	Nov 05, 2015
<u>AB</u>	TEVA PHARMS USA	<u>500MG;EQ 15MG BASE</u>	<u>A091155</u>	<u>001</u>	Mar 10, 2014
<u>AB</u>		<u>850MG;EQ 15MG BASE</u>	<u>A091155</u>	<u>002</u>	Mar 10, 2014

METFORMIN HYDROCHLORIDE; SAXAGLIPTIN HYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL

METFORMIN HYDROCHLORIDE; SAXAGLIPTIN HYDROCHLORIDE

<u>AB</u>	DR REDDYS LABS SA	<u>500MG;EQ 5MG BASE</u>	<u>A207678</u>	<u>001</u>	Aug 09, 2023
<u>AB</u>		<u>1GM;EQ 2.5MG BASE</u>	<u>A207678</u>	<u>002</u>	Aug 09, 2023
<u>AB</u>	!	<u>1GM;EQ 5MG BASE</u>	<u>A207678</u>	<u>003</u>	Aug 09, 2023
<u>SAXAGLIPTIN AND METFORMIN HYDROCHLORIDE</u>					
<u>AB</u>	MYLAN	<u>500MG;EQ 5MG BASE</u>	<u>A205981</u>	<u>001</u>	Jul 31, 2023
<u>AB</u>		<u>1GM;EQ 2.5MG BASE</u>	<u>A205981</u>	<u>002</u>	Jul 31, 2023
<u>AB</u>		<u>1GM;EQ 5MG BASE</u>	<u>A205981</u>	<u>003</u>	Jul 31, 2023
<u>AB</u>	SUN PHARM	<u>500MG;EQ 5MG BASE</u>	<u>A206081</u>	<u>001</u>	Jul 31, 2023
<u>AB</u>		<u>1GM;EQ 2.5MG BASE</u>	<u>A206081</u>	<u>002</u>	Jul 31, 2023
<u>AB</u>		<u>1GM;EQ 5MG BASE</u>	<u>A206081</u>	<u>003</u>	Jul 31, 2023

PRESCRIPTION DRUG PRODUCT LIST

METFORMIN HYDROCHLORIDE; SITAGLIPTIN

TABLET; ORAL

ZITUVIMET

+	ZYDUS LIFESCIENCES	500MG;50MG	N216743	001	Nov 03, 2023
+	!	1GM;50MG	N216743	002	Nov 03, 2023

TABLET, EXTENDED RELEASE; ORAL

ZITUVIMET XR

+	ZYDUS LIFESCIENCES	500MG;50MG	N216778	001	Jul 18, 2024
+		1GM;50MG	N216778	002	Jul 18, 2024
+	!	1GM;100MG	N216778	003	Jul 18, 2024

METFORMIN HYDROCHLORIDE; SITAGLIPTIN PHOSPHATE

TABLET; ORAL

JANUMET

+	MSD SUB MERCK	500MG;EQ 50MG BASE	N022044	001	Mar 30, 2007
+	!	1GM;EQ 50MG BASE	N022044	002	Mar 30, 2007

TABLET, EXTENDED RELEASE; ORAL

JANUMET XR

+	MSD SUB MERCK	500MG;EQ 50MG BASE	N202270	001	Feb 02, 2012
+		1GM;EQ 50MG BASE	N202270	002	Feb 02, 2012
+	!	1GM;EQ 100MG BASE	N202270	003	Feb 02, 2012

METHACHOLINE CHLORIDE

FOR SOLUTION; INHALATION

PROVOCHOLINE

+	METHAPHARM	100MG/VIAL	N019193	001	Oct 31, 1986
---	------------	------------	---------	-----	--------------

METHADONE HYDROCHLORIDE

CONCENTRATE; ORAL

METHADONE HYDROCHLORIDE

<u>AA</u>		HIKMA	<u>10MG/ML</u>	<u>A040180</u>	<u>001</u>	Apr 30, 1998
<u>AA</u>		LANNETT CO INC	<u>10MG/ML</u>	<u>A212093</u>	<u>001</u>	Nov 02, 2020
<u>AA</u>		SPECGX LLC	<u>10MG/ML</u>	<u>A207368</u>	<u>001</u>	Aug 22, 2019
<u>AA</u>		VISTAPHARM LLC	<u>10MG/ML</u>	<u>A040088</u>	<u>001</u>	Nov 30, 1994

METHADONE HYDROCHLORIDE INTENSOL

<u>AA</u>		HIKMA	<u>10MG/ML</u>	<u>A089897</u>	<u>001</u>	Sep 06, 1988
-----------	--	-------	----------------	----------------	------------	--------------

METHADOSE

<u>AA</u>	+	!	SPECGX LLC	<u>10MG/ML</u>	<u>N017116</u>	<u>002</u>
-----------	---	---	------------	----------------	----------------	------------

INJECTABLE; INJECTION

METHADONE HYDROCHLORIDE

<u>AP</u>		BRECKENRIDGE	<u>10MG/ML</u>	<u>A218252</u>	<u>001</u>	Dec 16, 2024
<u>AP</u>		LONG GROVE PHARMS	<u>10MG/ML</u>	<u>A208306</u>	<u>001</u>	Oct 27, 2017
<u>AP</u>	+	!	MYLAN INSTITUTIONAL	<u>10MG/ML</u>	<u>N021624</u>	<u>001</u>

SOLUTION; ORAL

METHADONE HYDROCHLORIDE

<u>AA</u>	+	!	HIKMA	<u>5MG/5ML</u>	<u>A087393</u>	<u>001</u>	
<u>AA</u>	+	!		<u>10MG/5ML</u>	<u>A087997</u>	<u>001</u>	Aug 30, 1982
<u>AA</u>			SPECGX LLC	<u>5MG/5ML</u>	<u>A207537</u>	<u>001</u>	Oct 02, 2019
<u>AA</u>				<u>10MG/5ML</u>	<u>A207537</u>	<u>002</u>	Oct 01, 2019

TABLET; ORAL

METHADONE HYDROCHLORIDE

<u>AA</u>		ASCENT PHARMS INC	<u>5MG</u>	<u>A211228</u>	<u>001</u>	Jan 03, 2019
<u>AA</u>			<u>10MG</u>	<u>A211228</u>	<u>002</u>	Jan 03, 2019
<u>AA</u>		AUROLIFE PHARMA LLC	<u>5MG</u>	<u>A203502</u>	<u>001</u>	Aug 31, 2015
<u>AA</u>			<u>10MG</u>	<u>A203502</u>	<u>002</u>	Aug 31, 2015
<u>AA</u>		ELITE LABS INC	<u>5MG</u>	<u>A210484</u>	<u>001</u>	Aug 02, 2018
<u>AA</u>			<u>10MG</u>	<u>A210484</u>	<u>002</u>	Aug 02, 2018
<u>AA</u>		EPIC PHARMA LLC	<u>5MG</u>	<u>A090065</u>	<u>001</u>	Aug 18, 2015
<u>AA</u>			<u>10MG</u>	<u>A090065</u>	<u>002</u>	Aug 18, 2015
<u>AA</u>		HIKMA	<u>5MG</u>	<u>A088108</u>	<u>001</u>	Mar 08, 1983
<u>AA</u>			<u>10MG</u>	<u>A088109</u>	<u>001</u>	Mar 08, 1983
<u>AA</u>	!		<u>5MG</u>	<u>A040517</u>	<u>001</u>	Apr 27, 2004
<u>AA</u>	!		<u>10MG</u>	<u>A040517</u>	<u>002</u>	Apr 27, 2004
<u>AA</u>		THEPHARMANETWORK LLC	<u>5MG</u>	<u>A090635</u>	<u>002</u>	Sep 22, 2020
<u>AA</u>			<u>10MG</u>	<u>A090635</u>	<u>001</u>	Nov 25, 2009
<u>AA</u>		VISTAPHARM LLC	<u>10MG</u>	<u>A040241</u>	<u>002</u>	May 29, 1998
<u>AA</u>			<u>10MG</u>	<u>A204166</u>	<u>001</u>	Mar 16, 2020

TABLET, FOR SUSPENSION; ORAL

METHADONE HYDROCHLORIDE

<u>AA</u>	+	!	HIKMA	<u>40MG</u>	<u>N017058</u>	<u>001</u>	
<u>AA</u>			SPECGX LLC	<u>40MG</u>	<u>A077142</u>	<u>001</u>	Jul 12, 2005
<u>AA</u>			VISTAPHARM	<u>40MG</u>	<u>A075082</u>	<u>001</u>	Mar 25, 1998

PRESCRIPTION DRUG PRODUCT LIST

METHADONE HYDROCHLORIDE

TABLET, FOR SUSPENSION;ORAL

METHADOSE

<u>AA</u>	SPECGX LLC	<u>40MG</u>	<u>A074184</u>	<u>001</u>	Apr 29, 1993
-----------	------------	-------------	----------------	------------	--------------

METHAMPHETAMINE HYDROCHLORIDE

TABLET;ORAL

DESOXYN

<u>AA</u>	+!	AJENAT PHARMS	<u>5MG</u>	<u>N005378</u>	<u>002</u>
-----------	----	---------------	------------	----------------	------------

METHAMPHETAMINE HYDROCHLORIDE

<u>AA</u>		DR REDDYS LABS SA	<u>5MG</u>	<u>A091189</u>	<u>001</u>	Apr 21, 2010
-----------	--	-------------------	------------	----------------	------------	--------------

<u>AA</u>		HIKMA	<u>5MG</u>	<u>A203846</u>	<u>001</u>	Nov 17, 2015
-----------	--	-------	------------	----------------	------------	--------------

METHAZOLAMIDE

TABLET;ORAL

METHAZOLAMIDE

<u>AB</u>		ANI PHARMS	<u>25MG</u>	<u>A040001</u>	<u>001</u>	Jun 30, 1993
-----------	--	------------	-------------	----------------	------------	--------------

<u>AB</u>			<u>50MG</u>	<u>A040001</u>	<u>002</u>	Jun 30, 1993
-----------	--	--	-------------	----------------	------------	--------------

<u>AB</u>		BAUSCH AND LOMB INC	<u>25MG</u>	<u>A207438</u>	<u>001</u>	Oct 05, 2018
-----------	--	---------------------	-------------	----------------	------------	--------------

<u>AB</u>			<u>50MG</u>	<u>A207438</u>	<u>002</u>	Oct 05, 2018
-----------	--	--	-------------	----------------	------------	--------------

<u>AB</u>		CHARTWELL RX	<u>25MG</u>	<u>A040036</u>	<u>001</u>	Jun 30, 1993
-----------	--	--------------	-------------	----------------	------------	--------------

<u>AB</u>			<u>50MG</u>	<u>A040036</u>	<u>002</u>	Jun 30, 1993
-----------	--	--	-------------	----------------	------------	--------------

<u>AB</u>		MIKART	<u>25MG</u>	<u>A040062</u>	<u>001</u>	Jan 27, 1994
-----------	--	--------	-------------	----------------	------------	--------------

<u>AB</u>	!		<u>50MG</u>	<u>A040062</u>	<u>002</u>	Jan 27, 1994
-----------	---	--	-------------	----------------	------------	--------------

<u>AB</u>		TAGI	<u>25MG</u>	<u>A215615</u>	<u>001</u>	Oct 18, 2022
-----------	--	------	-------------	----------------	------------	--------------

<u>AB</u>			<u>50MG</u>	<u>A215615</u>	<u>002</u>	Oct 18, 2022
-----------	--	--	-------------	----------------	------------	--------------

METHENAMINE HIPPURATE

TABLET;ORAL

HIPREX

<u>AB</u>	+	ESJAY PHARMA	<u>1GM</u>	<u>N017681</u>	<u>001</u>
-----------	---	--------------	------------	----------------	------------

METHENAMINE HIPPURATE

<u>AB</u>	!	AUROBINDO PHARMA LTD	<u>1GM</u>	<u>A205661</u>	<u>001</u>	Jul 05, 2016
-----------	---	----------------------	------------	----------------	------------	--------------

<u>AB</u>		JUBILANT CADISTA	<u>1GM</u>	<u>A217675</u>	<u>001</u>	Dec 01, 2023
-----------	--	------------------	------------	----------------	------------	--------------

<u>AB</u>		MICRO LABS	<u>1GM</u>	<u>A212172</u>	<u>001</u>	Aug 01, 2019
-----------	--	------------	------------	----------------	------------	--------------

<u>AB</u>		NOVAST LABS	<u>1GM</u>	<u>A210068</u>	<u>001</u>	Nov 27, 2020
-----------	--	-------------	------------	----------------	------------	--------------

UREX

<u>AB</u>		ALVOGEN	<u>1GM</u>	<u>N016151</u>	<u>001</u>
-----------	--	---------	------------	----------------	------------

METHIMAZOLE

TABLET;ORAL

METHIMAZOLE

<u>AB</u>		BIONPHARMA	<u>5MG</u>	<u>A218149</u>	<u>001</u>	Sep 25, 2023
-----------	--	------------	------------	----------------	------------	--------------

<u>AB</u>			<u>10MG</u>	<u>A218149</u>	<u>002</u>	Sep 25, 2023
-----------	--	--	-------------	----------------	------------	--------------

<u>AB</u>		CHARTWELL RX	<u>5MG</u>	<u>A040411</u>	<u>001</u>	Mar 27, 2001
-----------	--	--------------	------------	----------------	------------	--------------

<u>AB</u>			<u>10MG</u>	<u>A040411</u>	<u>002</u>	Mar 27, 2001
-----------	--	--	-------------	----------------	------------	--------------

<u>AB</u>		HERITAGE PHARMA	<u>5MG</u>	<u>A040734</u>	<u>001</u>	Dec 14, 2007
-----------	--	-----------------	------------	----------------	------------	--------------

<u>AB</u>			<u>10MG</u>	<u>A040734</u>	<u>002</u>	Dec 14, 2007
-----------	--	--	-------------	----------------	------------	--------------

<u>AB</u>		MACLEODS PHARMS LTD	<u>5MG</u>	<u>A209827</u>	<u>001</u>	May 24, 2023
-----------	--	---------------------	------------	----------------	------------	--------------

<u>AB</u>			<u>10MG</u>	<u>A209827</u>	<u>002</u>	May 24, 2023
-----------	--	--	-------------	----------------	------------	--------------

<u>AB</u>		MYLAN	<u>5MG</u>	<u>A040350</u>	<u>001</u>	Mar 29, 2000
-----------	--	-------	------------	----------------	------------	--------------

<u>AB</u>	!		<u>10MG</u>	<u>A040350</u>	<u>002</u>	Mar 29, 2000
-----------	---	--	-------------	----------------	------------	--------------

<u>AB</u>		QINGDAO BAHEAL PHARM	<u>5MG</u>	<u>A040547</u>	<u>001</u>	Feb 18, 2005
-----------	--	----------------------	------------	----------------	------------	--------------

<u>AB</u>			<u>10MG</u>	<u>A040547</u>	<u>002</u>	Feb 18, 2005
-----------	--	--	-------------	----------------	------------	--------------

<u>AB</u>		RISING	<u>5MG</u>	<u>A202068</u>	<u>001</u>	Mar 07, 2012
-----------	--	--------	------------	----------------	------------	--------------

<u>AB</u>			<u>10MG</u>	<u>A202068</u>	<u>002</u>	Mar 07, 2012
-----------	--	--	-------------	----------------	------------	--------------

METHOCARBAMOL

SOLUTION;IM-IV

METHOCARBAMOL

<u>AP</u>		AM REGENT	<u>1GM/10ML (100MG/ML)</u>	<u>A207496</u>	<u>001</u>	Jun 22, 2017
-----------	--	-----------	----------------------------	----------------	------------	--------------

<u>AP</u>		EUGIA PHARMA	<u>1GM/10ML (100MG/ML)</u>	<u>A206128</u>	<u>001</u>	May 27, 2016
-----------	--	--------------	----------------------------	----------------	------------	--------------

<u>AP</u>		FRESENIUS KABI USA	<u>1GM/10ML (100MG/ML)</u>	<u>A209331</u>	<u>001</u>	Apr 17, 2018
-----------	--	--------------------	----------------------------	----------------	------------	--------------

<u>AP</u>		GLAND PHARMA LTD	<u>1GM/10ML (100MG/ML)</u>	<u>A211504</u>	<u>001</u>	Oct 26, 2018
-----------	--	------------------	----------------------------	----------------	------------	--------------

<u>AP</u>		MONTEREY PHARMS LLC	<u>1GM/10ML (100MG/ML)</u>	<u>A205354</u>	<u>001</u>	Oct 27, 2016
-----------	--	---------------------	----------------------------	----------------	------------	--------------

<u>AP</u>		SAGENT PHARMS INC	<u>1GM/10ML (100MG/ML)</u>	<u>A205404</u>	<u>001</u>	Jul 18, 2017
-----------	--	-------------------	----------------------------	----------------	------------	--------------

<u>AP</u>		SLATE RUN PHARMA	<u>1GM/10ML (100MG/ML)</u>	<u>A208116</u>	<u>001</u>	Jan 19, 2017
-----------	--	------------------	----------------------------	----------------	------------	--------------

<u>AP</u>		SOMERSET THERAPS LLC	<u>1GM/10ML (100MG/ML)</u>	<u>A207522</u>	<u>001</u>	Jul 31, 2017
-----------	--	----------------------	----------------------------	----------------	------------	--------------

ROBAXIN

<u>AP</u>	+	HIKMA	<u>1GM/10ML (100MG/ML)</u>	<u>N011790</u>	<u>001</u>
-----------	---	-------	----------------------------	----------------	------------

PRESCRIPTION DRUG PRODUCT LIST

METHOCARBAMOL

TABLET; ORAL

METHOCARBAMOL

<u>AA</u>	ALLIED	<u>500MG</u>	<u>A212623</u>	<u>001</u>	Apr 30, 2021
<u>AA</u>		<u>750MG</u>	<u>A212623</u>	<u>002</u>	Apr 30, 2021
<u>AA</u>	AUROBINDO PHARMA LTD	<u>500MG</u>	<u>A213967</u>	<u>002</u>	Feb 09, 2023
<u>AA</u>		<u>750MG</u>	<u>A213967</u>	<u>001</u>	Aug 12, 2020
<u>AA</u>	BEXIMCO PHARMS USA	<u>500MG</u>	<u>A208507</u>	<u>001</u>	Jul 21, 2017
<u>AA</u>		<u>750MG</u>	<u>A208507</u>	<u>002</u>	Jul 21, 2017
<u>AA</u>	!	<u>500MG</u>	<u>A209312</u>	<u>001</u>	May 07, 2018
<u>AA</u>	!	<u>750MG</u>	<u>A209312</u>	<u>002</u>	May 07, 2018
<u>AA</u>	HANGZHOU BINJIANG	<u>500MG</u>	<u>A200958</u>	<u>001</u>	Oct 21, 2011
<u>AA</u>		<u>750MG</u>	<u>A200958</u>	<u>002</u>	Oct 21, 2011
<u>AA</u>	HETERO LABS LTD III	<u>500MG</u>	<u>A090200</u>	<u>001</u>	Nov 06, 2009
<u>AA</u>		<u>750MG</u>	<u>A090200</u>	<u>002</u>	Nov 06, 2009
<u>AA</u>	HLTHCARE	<u>500MG</u>	<u>A203550</u>	<u>001</u>	Feb 08, 2017
<u>AA</u>		<u>750MG</u>	<u>A203550</u>	<u>002</u>	Feb 08, 2017
<u>AA</u>	OXFORD PHARMS	<u>500MG</u>	<u>A040489</u>	<u>001</u>	Jan 29, 2003
<u>AA</u>		<u>750MG</u>	<u>A040489</u>	<u>002</u>	Jan 29, 2003
<u>AA</u>	PRINSTON INC	<u>500MG</u>	<u>A086988</u>	<u>002</u>	
<u>AA</u>		<u>750MG</u>	<u>A086988</u>	<u>001</u>	
<u>AB</u>	!	<u>1GM</u>	<u>A200958</u>	<u>003</u>	Dec 06, 2021
<u>AB</u>	MIKART	<u>1GM</u>	<u>A212707</u>	<u>001</u>	Jun 12, 2023

METHOHEXITAL SODIUM

INJECTABLE; INJECTION

BREVITAL SODIUM

<u>AP</u>	+!	ENDO OPERATIONS	<u>500MG/VIAL</u>	<u>N011559</u>	<u>001</u>
-----------	----	-----------------	-------------------	----------------	------------

METHOHEXITAL SODIUM

<u>AP</u>		STERISCIENCE SPECLTS	<u>500MG/VIAL</u>	<u>A215488</u>	<u>001</u>	Oct 03, 2024
-----------	--	-------------------------	-------------------	----------------	------------	--------------

METHOTREXATE

SOLUTION; ORAL

JYLAMVO

+!	SHORLA	2MG/ML	N212479	001	Nov 29, 2022
----	--------	--------	---------	-----	--------------

SOLUTION; SUBCUTANEOUS

OTREXUP

+!	OTTER PHARMS	10MG/0.4ML (10MG/0.4ML)	N204824	001	Oct 11, 2013
+!		12.5MG/0.4ML (12.5MG/0.4ML)	N204824	006	Mar 24, 2016
+!		15MG/0.4ML (15MG/0.4ML)	N204824	002	Oct 11, 2013
+!		17.5MG/0.4ML (17.5MG/0.4ML)	N204824	007	Mar 24, 2016
+!		20MG/0.4ML (20MG/0.4ML)	N204824	003	Oct 11, 2013
+!		22.5MG/0.4ML (22.5MG/0.4ML)	N204824	008	Mar 24, 2016
+!		25MG/0.4ML (25MG/0.4ML)	N204824	004	Oct 11, 2013

RASUVO

+!	MEDEXUS	7.5MG/0.15ML (7.5MG/0.15ML)	N205776	001	Jul 10, 2014
+!		10MG/0.20ML (10MG/0.20ML)	N205776	002	Jul 10, 2014
+!		12.5MG/0.25ML (12.5MG/0.25ML)	N205776	003	Jul 10, 2014
+!		15MG/0.30ML (15MG/0.30ML)	N205776	004	Jul 10, 2014
+!		17.5MG/0.35ML (17.5MG/0.35ML)	N205776	005	Jul 10, 2014
+!		20MG/0.4ML (20MG/0.4ML)	N205776	006	Jul 10, 2014
+!		22.5MG/0.45ML (22.5MG/0.45ML)	N205776	007	Jul 10, 2014
+!		25MG/0.5ML (25MG/0.5ML)	N205776	008	Jul 10, 2014
+!		30MG/0.6ML (30MG/0.6ML)	N205776	010	Jul 10, 2014

METHOTREXATE SODIUM

INJECTABLE; INJECTION

METHOTREXATE PRESERVATIVE FREE

<u>AP</u>		FRESENIUS KABI USA	<u>EQ 1GM BASE/VIAL</u>	<u>A040266</u>	<u>001</u>	Feb 26, 1999
-----------	--	--------------------	-------------------------	----------------	------------	--------------

METHOTREXATE SODIUM

<u>AP</u>	!	FRESENIUS KABI USA	<u>EQ 50MG BASE/2ML (EQ 25MG BASE/ML)</u>	<u>A040263</u>	<u>001</u>	Feb 26, 1999
<u>AP</u>	!	HIKMA	<u>EQ 100MG BASE/4ML (EQ 25MG BASE/ML)</u>	<u>A089341</u>	<u>001</u>	Sep 16, 1986
<u>AP</u>	+!	HOSPIRA	<u>EQ 50MG BASE/2ML (EQ 25MG BASE/ML)</u>	<u>N011719</u>	<u>010</u>	Dec 15, 2004

METHOTREXATE SODIUM PRESERVATIVE FREE

<u>AP</u>	!	ACCORD HLTHCARE	<u>EQ 50MG BASE/2ML (EQ 25MG BASE/ML)</u>	<u>A040767</u>	<u>001</u>	Apr 30, 2007
<u>AP</u>	!		<u>EQ 250MG BASE/10ML (EQ 25MG BASE/ML)</u>	<u>A040768</u>	<u>001</u>	Apr 30, 2007
<u>AP</u>	!		<u>EQ 1GM BASE/40ML (EQ 25MG BASE/ML)</u>	<u>A040716</u>	<u>001</u>	Apr 30, 2007
<u>AP</u>	!	HIKMA	<u>EQ 50MG BASE/2ML (EQ 25MG BASE/ML)</u>	<u>A089340</u>	<u>001</u>	Sep 16, 1986
<u>AP</u>			<u>EQ 250MG BASE/10ML (EQ 25MG BASE/ML)</u>	<u>A089343</u>	<u>001</u>	Sep 16, 1986
<u>AP</u>	!		<u>EQ 1GM BASE/VIAL</u>	<u>A040632</u>	<u>001</u>	Aug 12, 2005
<u>AP</u>	+!	HOSPIRA	<u>EQ 1GM BASE/40ML (EQ 25MG BASE/ML)</u>	<u>N011719</u>	<u>012</u>	Apr 13, 2005
<u>AP</u>		PHARMACHEMIE BV	<u>EQ 50MG BASE/2ML (EQ 25MG BASE/ML)</u>	<u>A040843</u>	<u>002</u>	Jan 11, 2010

PRESCRIPTION DRUG PRODUCT LIST

METHOTREXATE SODIUM

INJECTABLE; INJECTION

METHOTREXATE SODIUM PRESERVATIVE FREE

<u>AP</u>		<u>EQ 250MG BASE/10ML (EQ 25MG BASE/ML)</u>	<u>A040843 004</u>	Jan 11, 2010
<u>AP</u>		<u>EQ 1GM BASE/40ML (EQ 25MG BASE/ML)</u>	<u>A040843 001</u>	Jan 11, 2010
<u>AP</u>	SAGENT PHARMS INC	<u>EQ 50MG BASE/2ML (EQ 25MG BASE/ML)</u>	<u>A203407 001</u>	Aug 09, 2018
<u>AP</u>		<u>EQ 250MG BASE/10ML (EQ 25MG BASE/ML)</u>	<u>A203407 002</u>	Aug 09, 2018
<u>AP</u>		<u>EQ 1GM BASE/40ML (EQ 25MG BASE/ML)</u>	<u>A203407 003</u>	Aug 09, 2018
<u>AP</u>	SANDOZ	<u>EQ 50MG BASE/2ML (EQ 25MG BASE/ML)</u>	<u>A090039 001</u>	Mar 31, 2009
<u>AP</u>		<u>EQ 250MG BASE/10ML (EQ 25MG BASE/ML)</u>	<u>A090039 002</u>	Mar 31, 2009
<u>AP</u>		<u>EQ 1GM BASE/40ML (EQ 25MG BASE/ML)</u>	<u>A090029 001</u>	Mar 31, 2009

METHOTREXATE SODIUM

! HIKMA

EQ 200MG BASE/8ML (EQ 25MG BASE/ML)

A089342 001 Sep 16, 1986

METHOTREXATE SODIUM PRESERVATIVE FREE

PHARMACHEMIE BV

EQ 100MG BASE/4ML (EQ 25MG BASE/ML)

A040843 003 Feb 27, 2012

SOLUTION; ORAL

XATMEP

+! AZURITY

EQ 2.5MG BASE/ML

N208400 001 Apr 25, 2017

TABLET; ORAL

METHOTREXATE SODIUM

<u>AB</u>	ACCORD HLTHCARE	<u>EQ 2.5MG BASE</u>	<u>A213343 001</u>	Jan 24, 2020
<u>AB</u>	ALEMBIC	<u>EQ 2.5MG BASE</u>	<u>A217552 001</u>	Jun 03, 2024
<u>AB</u>	BARR	<u>EQ 2.5MG BASE</u>	<u>A081099 001</u>	Oct 15, 1990
<u>AB</u>	DAITO	<u>EQ 2.5MG BASE</u>	<u>A213362 001</u>	Aug 07, 2023
<u>AB</u>	ELITE LABS INC	<u>EQ 2.5MG BASE</u>	<u>A216453 001</u>	May 16, 2024
<u>AB</u>	EUGIA PHARMA	<u>EQ 2.5MG BASE</u>	<u>A210454 001</u>	Jan 30, 2020
<u>AB</u>	! HIKMA	<u>EQ 2.5MG BASE</u>	<u>A040054 001</u>	Aug 01, 1994
<u>AB</u>	MYLAN	<u>EQ 2.5MG BASE</u>	<u>A081235 001</u>	May 15, 1992
<u>AB</u>	SUN PHARM	<u>EQ 2.5MG BASE</u>	<u>A201749 001</u>	May 21, 2015
<u>AB</u>	ZYDUS PHARMS	<u>EQ 2.5MG BASE</u>	<u>A207812 001</u>	Jan 13, 2017
	TREXALL			
	BARR	EQ 5MG BASE	A040385 001	Mar 21, 2001
		EQ 7.5MG BASE	A040385 002	Mar 21, 2001
		EQ 10MG BASE	A040385 003	Mar 21, 2001
	!	EQ 15MG BASE	A040385 004	Mar 21, 2001

METHOXSALLEN

CAPSULE; ORAL

METHOXSALLEN

<u>AB</u>	STRIDES SOFTGELS	<u>10MG</u>	<u>A202687 001</u>	Jun 05, 2014
	<u>OXSORALEN-ULTRA</u>			
<u>AB</u>	+! BAUSCH	<u>10MG</u>	<u>N019600 001</u>	Oct 30, 1986
	INJECTABLE; INJECTION			
	UVADEX			
	+! THERAKOS	0.02MG/ML	N020969 001	Feb 25, 1999
	DEVELOPMENT			

METHSCOPOLAMINE BROMIDE

TABLET; ORAL

METHSCOPOLAMINE BROMIDE

<u>AA</u>	BRECKENRIDGE PHARM	<u>2.5MG</u>	<u>A040642 001</u>	Dec 06, 2011
<u>AA</u>		<u>5MG</u>	<u>A040642 002</u>	Dec 06, 2011
<u>AA</u>	NE RX PHARMA	<u>2.5MG</u>	<u>A216786 001</u>	Feb 01, 2024
<u>AA</u>		<u>5MG</u>	<u>A216786 002</u>	Feb 01, 2024
<u>AA</u>	UNICHEM	<u>2.5MG</u>	<u>A200602 001</u>	Sep 24, 2012
<u>AA</u>	!	<u>5MG</u>	<u>A200602 002</u>	Sep 24, 2012

METHSUXIMIDE

CAPSULE; ORAL

CELONTIN

<u>AB</u>	+! PARKE DAVIS	<u>300MG</u>	<u>N010596 008</u>	
-----------	----------------	--------------	--------------------	--

METHSUXIMIDE

<u>AB</u>	NOVITIUM PHARMA	<u>300MG</u>	<u>A217213 001</u>	May 01, 2023
-----------	-----------------	--------------	--------------------	--------------

METHYLDOPA

TABLET; ORAL

METHYLDOPA

! RISING

250MG

A070076 002 Apr 18, 1985

!

500MG

A070076 001 Apr 18, 1985

PRESCRIPTION DRUG PRODUCT LIST

METHYLENE BLUE

SOLUTION; INTRAVENOUS

METHYLENE BLUE

<u>AP</u>	NEXUS	<u>50MG/10ML (5MG/ML)</u>	<u>A217561 001</u>	Nov 21, 2024
<u>AP</u>	RK PHARMA	<u>50MG/10ML (5MG/ML)</u>	<u>A216959 001</u>	Dec 17, 2024
<u>AP</u>	STERISCIENCE	<u>50MG/10ML (5MG/ML)</u>	<u>A216955 001</u>	Nov 22, 2024
<u>AP</u>	ZYDUS LIFESCIENCES	<u>10MG/2ML (5MG/ML)</u>	<u>A215636 001</u>	Dec 05, 2023
<u>AP</u>		<u>50MG/10ML (5MG/ML)</u>	<u>A215636 002</u>	Dec 05, 2023
<u>PROVAYBLUE</u>				
<u>AP</u>	+! PROVEPHARM SAS	<u>10MG/2ML (5MG/ML)</u>	<u>N204630 002</u>	Jul 18, 2019
<u>AP</u>	+!	<u>50MG/10ML (5MG/ML)</u>	<u>N204630 001</u>	Apr 08, 2016

METHYLERGONOVINE MALEATE

INJECTABLE; INJECTION

METHYLERGONOVINE MALEATE

<u>AP</u>	! AM REGENT	<u>0.2MG/ML</u>	<u>A090193 001</u>	Nov 24, 2008
<u>AP</u>	BRECKENRIDGE	<u>0.2MG/ML</u>	<u>A040889 001</u>	Sep 13, 2010

TABLET; ORAL

METHYLERGONOVINE MALEATE

<u>AB</u>	AMNEAL PHARMS	<u>0.2MG</u>	<u>A211483 001</u>	Sep 10, 2018
<u>AB</u>	! CHARTWELL RX	<u>0.2MG</u>	<u>A091577 001</u>	May 02, 2011
<u>AB</u>	GRANULES	<u>0.2MG</u>	<u>A210424 001</u>	May 15, 2018
<u>AB</u>	RISING	<u>0.2MG</u>	<u>A211919 001</u>	Jan 15, 2021
<u>AB</u>	TEVA PHARMS USA	<u>0.2MG</u>	<u>A211455 001</u>	Mar 20, 2019
<u>AB</u>	TRUPHARMA	<u>0.2MG</u>	<u>A212233 001</u>	May 01, 2020

METHYLNALTREXONE BROMIDE

SOLUTION; SUBCUTANEOUS

METHYLNALTREXONE BROMIDE

<u>AP</u>	ACTAVIS LLC	<u>8MG/0.4ML (8MG/0.4ML)</u>	<u>A208112 001</u>	Aug 26, 2024
<u>AP</u>		<u>12MG/0.6ML (12MG/0.6ML)</u>	<u>A208112 002</u>	Aug 26, 2024

RELISTOR

<u>AP</u>	+! SALIX PHARMS	<u>8MG/0.4ML (8MG/0.4ML)</u>	<u>N021964 002</u>	Sep 27, 2010
<u>AP</u>	+!	<u>12MG/0.6ML (12MG/0.6ML)</u>	<u>N021964 003</u>	Sep 27, 2010
	+!	12MG/0.6ML (12MG/0.6ML)	N021964 001	Apr 24, 2008

TABLET; ORAL

RELISTOR

+!	SALIX	150MG	N208271 001	Jul 19, 2016
----	-------	-------	-------------	--------------

METHYLPHENIDATE

FILM, EXTENDED RELEASE; TRANSDERMAL

DAYTRANA

<u>AB</u>	+ NOVEN PHARMS INC	<u>10MG/9HR (1.1MG/HR)</u>	<u>N021514 001</u>	Apr 06, 2006
<u>AB</u>	+	<u>15MG/9HR (1.6MG/HR)</u>	<u>N021514 002</u>	Apr 06, 2006
<u>AB</u>	+	<u>20MG/9HR (2.2MG/HR)</u>	<u>N021514 003</u>	Apr 06, 2006
<u>AB</u>	+!	<u>30MG/9HR (3.3MG/HR)</u>	<u>N021514 004</u>	Apr 06, 2006

METHYLPHENIDATE

<u>AB</u>	MYLAN TECH VIATRIS	<u>10MG/9HR (1.1MG/HR)</u>	<u>A206497 001</u>	Mar 14, 2022
<u>AB</u>		<u>15MG/9HR (1.6MG/HR)</u>	<u>A206497 002</u>	Mar 14, 2022
<u>AB</u>		<u>20MG/9HR (2.2MG/HR)</u>	<u>A206497 003</u>	Mar 14, 2022
<u>AB</u>		<u>30MG/9HR (3.3MG/HR)</u>	<u>A206497 004</u>	Mar 14, 2022

TABLET, ORALLY DISINTEGRATING, EXTENDED RELEASE; ORAL

COTEMPLA XR-ODT

+	NEOS THERAPS INC	8.6MG	N205489 001	Jun 19, 2017
+		17.3MG	N205489 002	Jun 19, 2017
+	!	25.9MG	N205489 003	Jun 19, 2017

METHYLPHENIDATE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

METHYLPHENIDATE HYDROCHLORIDE

<u>AB1</u>	DR REDDYS LABS SA	<u>10MG</u>	<u>A200886 001</u>	Feb 26, 2018
<u>AB1</u>		<u>20MG</u>	<u>A078458 001</u>	Dec 01, 2011
<u>AB1</u>		<u>30MG</u>	<u>A078458 002</u>	Dec 01, 2011
<u>AB1</u>		<u>40MG</u>	<u>A078458 003</u>	Dec 01, 2011
<u>AB1</u>	!	<u>60MG</u>	<u>A078458 004</u>	Jun 23, 2016
<u>AB1</u>	GRANULES	<u>10MG</u>	<u>A211796 001</u>	May 23, 2019
<u>AB1</u>		<u>20MG</u>	<u>A211796 002</u>	May 23, 2019
<u>AB1</u>		<u>30MG</u>	<u>A211796 003</u>	May 23, 2019
<u>AB1</u>		<u>40MG</u>	<u>A211796 004</u>	May 23, 2019
<u>AB1</u>		<u>60MG</u>	<u>A211796 005</u>	May 23, 2019

RITALIN LA

<u>AB1</u>	+ SANDOZ	<u>10MG</u>	<u>N021284 004</u>	Apr 10, 2004
<u>AB1</u>	+	<u>20MG</u>	<u>N021284 001</u>	Jun 05, 2002
<u>AB1</u>	+	<u>30MG</u>	<u>N021284 002</u>	Jun 05, 2002

PRESCRIPTION DRUG PRODUCT LIST

METHYLPHENIDATE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE;ORAL

RITALIN LA

<u>AB1</u>	+		<u>40MG</u>	<u>N021284</u>	<u>003</u>	Jun 05, 2002
------------	---	--	-------------	----------------	------------	--------------

METADATE CD

<u>AB2</u>	+	AYTU BIOPHARMA	<u>10MG</u>	<u>N021259</u>	<u>003</u>	May 27, 2003
<u>AB2</u>	+		<u>20MG</u>	<u>N021259</u>	<u>001</u>	Apr 03, 2001
<u>AB2</u>	+		<u>30MG</u>	<u>N021259</u>	<u>002</u>	Jun 19, 2003
<u>AB2</u>	+		<u>40MG</u>	<u>N021259</u>	<u>004</u>	Feb 19, 2006
<u>AB2</u>	+		<u>50MG</u>	<u>N021259</u>	<u>005</u>	Feb 19, 2006
<u>AB2</u>	+		<u>60MG</u>	<u>N021259</u>	<u>006</u>	Feb 19, 2006

METHYLPHENIDATE HYDROCHLORIDE

<u>AB2</u>		IMPAX LABS INC	<u>10MG</u>	<u>A205105</u>	<u>001</u>	Jul 28, 2016
<u>AB2</u>			<u>20MG</u>	<u>A205105</u>	<u>002</u>	Jul 28, 2016
<u>AB2</u>			<u>30MG</u>	<u>A205105</u>	<u>003</u>	Jul 28, 2016
<u>AB2</u>			<u>40MG</u>	<u>A205105</u>	<u>004</u>	Jul 28, 2016
<u>AB2</u>			<u>50MG</u>	<u>A205105</u>	<u>005</u>	Jul 28, 2016
<u>AB2</u>			<u>60MG</u>	<u>A205105</u>	<u>006</u>	Jul 28, 2016
<u>AB2</u>		SPECGX LLC	<u>10MG</u>	<u>A203583</u>	<u>001</u>	Sep 29, 2015
<u>AB2</u>			<u>20MG</u>	<u>A203583</u>	<u>002</u>	Sep 29, 2015
<u>AB2</u>			<u>30MG</u>	<u>A203583</u>	<u>003</u>	Sep 29, 2015
<u>AB2</u>			<u>40MG</u>	<u>A203583</u>	<u>004</u>	Sep 29, 2015
<u>AB2</u>			<u>50MG</u>	<u>A203583</u>	<u>005</u>	Sep 29, 2015
<u>AB2</u>			<u>60MG</u>	<u>A203583</u>	<u>006</u>	Sep 29, 2015
<u>AB2</u>		TEVA PHARMS	<u>10MG</u>	<u>A077707</u>	<u>001</u>	Jul 19, 2012
<u>AB2</u>			<u>20MG</u>	<u>A077707</u>	<u>002</u>	Jul 19, 2012
<u>AB2</u>			<u>30MG</u>	<u>A077707</u>	<u>003</u>	Jul 19, 2012

APTENSIO XR

<u>AB3</u>	+	RHODES PHARMS	<u>10MG</u>	<u>N205831</u>	<u>001</u>	Apr 17, 2015
<u>AB3</u>	+		<u>15MG</u>	<u>N205831</u>	<u>002</u>	Apr 17, 2015
<u>AB3</u>	+		<u>20MG</u>	<u>N205831</u>	<u>003</u>	Apr 17, 2015
<u>AB3</u>	+		<u>30MG</u>	<u>N205831</u>	<u>004</u>	Apr 17, 2015
<u>AB3</u>	+		<u>40MG</u>	<u>N205831</u>	<u>005</u>	Apr 17, 2015
<u>AB3</u>	+		<u>50MG</u>	<u>N205831</u>	<u>006</u>	Apr 17, 2015
<u>AB3</u>	+		<u>60MG</u>	<u>N205831</u>	<u>007</u>	Apr 17, 2015

METHYLPHENIDATE HYDROCHLORIDE

<u>AB3</u>		ACTAVIS ELIZABETH	<u>10MG</u>	<u>A208861</u>	<u>001</u>	Dec 13, 2018
<u>AB3</u>			<u>15MG</u>	<u>A208861</u>	<u>002</u>	Dec 13, 2018
<u>AB3</u>			<u>20MG</u>	<u>A208861</u>	<u>003</u>	Dec 13, 2018
<u>AB3</u>			<u>30MG</u>	<u>A208861</u>	<u>004</u>	Dec 13, 2018
<u>AB3</u>			<u>40MG</u>	<u>A208861</u>	<u>005</u>	Dec 13, 2018
<u>AB3</u>			<u>50MG</u>	<u>A208861</u>	<u>006</u>	Dec 13, 2018
<u>AB3</u>			<u>60MG</u>	<u>A208861</u>	<u>007</u>	Dec 13, 2018

JORNAY PM

	+	IRONSHORE PHARMS	20MG	N209311	001	Aug 08, 2018
	+		40MG	N209311	002	Aug 08, 2018
	+		60MG	N209311	003	Aug 08, 2018
	+		80MG	N209311	004	Aug 08, 2018
	+		100MG	N209311	005	Aug 08, 2018

FOR SUSPENSION, EXTENDED RELEASE;ORAL

METHYLPHENIDATE HYDROCHLORIDE

<u>AB</u>		ACTAVIS LABS FL INC	<u>5MG/ML</u>	<u>A206049</u>	<u>001</u>	May 17, 2018
-----------	--	---------------------	---------------	----------------	------------	--------------

QUILLIVANT XR

<u>AB</u>	+	NEXTWAVE	<u>5MG/ML</u>	<u>N202100</u>	<u>001</u>	Sep 27, 2012
-----------	---	----------	---------------	----------------	------------	--------------

SOLUTION;ORAL

METHYLIN

<u>AA</u>	+	SPECGX LLC	<u>5MG/5ML</u>	<u>N021419</u>	<u>001</u>	Dec 19, 2002
<u>AA</u>	+		<u>10MG/5ML</u>	<u>N021419</u>	<u>002</u>	Dec 19, 2002

METHYLPHENIDATE HYDROCHLORIDE

<u>AA</u>		ABHAI LLC	<u>5MG/5ML</u>	<u>A207485</u>	<u>001</u>	Nov 18, 2016
<u>AA</u>			<u>10MG/5ML</u>	<u>A207485</u>	<u>002</u>	Nov 18, 2016
<u>AA</u>		ALKEM LABS LTD	<u>5MG/5ML</u>	<u>A211647</u>	<u>001</u>	Mar 30, 2020
<u>AA</u>			<u>10MG/5ML</u>	<u>A211647</u>	<u>002</u>	Mar 30, 2020
<u>AA</u>		ANDA REPOSITORY	<u>5MG/5ML</u>	<u>A210764</u>	<u>001</u>	Apr 10, 2020
<u>AA</u>			<u>10MG/5ML</u>	<u>A210764</u>	<u>002</u>	Apr 10, 2020
<u>AA</u>		ASCENT PHARMS INC	<u>5MG/5ML</u>	<u>A207417</u>	<u>001</u>	Jan 29, 2021
<u>AA</u>			<u>10MG/5ML</u>	<u>A207417</u>	<u>002</u>	Jan 29, 2021
<u>AA</u>		NOVEL LABS INC	<u>5MG/5ML</u>	<u>A204602</u>	<u>001</u>	Aug 14, 2015
<u>AA</u>			<u>10MG/5ML</u>	<u>A204602</u>	<u>002</u>	Aug 14, 2015
<u>AA</u>		QUAGEN	<u>5MG/5ML</u>	<u>A213567</u>	<u>001</u>	Jun 04, 2020
<u>AA</u>			<u>10MG/5ML</u>	<u>A213567</u>	<u>002</u>	Jun 04, 2020
<u>AA</u>		TRIS PHARMA INC	<u>5MG/5ML</u>	<u>A091601</u>	<u>001</u>	Jul 23, 2010

PRESCRIPTION DRUG PRODUCT LIST

METHYLPHENIDATE HYDROCHLORIDE

SOLUTION;ORAL

METHYLPHENIDATE HYDROCHLORIDE

<u>AA</u>		<u>10MG/5ML</u>	<u>A091601 002</u>	Jul 23, 2010
<u>AA</u>	WES PHARMA INC	<u>5MG/5ML</u>	<u>A210139 001</u>	Oct 03, 2018
<u>AA</u>		<u>10MG/5ML</u>	<u>A210139 002</u>	Oct 03, 2018

TABLET;ORAL

METHYLPHENIDATE HYDROCHLORIDE

<u>AB</u>	ABHAI INC	<u>5MG</u>	<u>A206932 001</u>	May 11, 2017
<u>AB</u>		<u>10MG</u>	<u>A206932 002</u>	May 11, 2017
<u>AB</u>		<u>20MG</u>	<u>A206932 003</u>	May 11, 2017
<u>AB</u>	ACCORD HLTHCARE	<u>5MG</u>	<u>A213936 001</u>	Oct 28, 2020
<u>AB</u>		<u>10MG</u>	<u>A213936 002</u>	Oct 28, 2020
<u>AB</u>		<u>20MG</u>	<u>A213936 003</u>	Oct 28, 2020
<u>AB</u>	ALKEM LABS LTD	<u>5MG</u>	<u>A211779 001</u>	Oct 04, 2019
<u>AB</u>		<u>10MG</u>	<u>A211779 002</u>	Oct 04, 2019
<u>AB</u>		<u>20MG</u>	<u>A211779 003</u>	Oct 04, 2019
<u>AB</u>	ASCENT PHARMS INC	<u>5MG</u>	<u>A207416 001</u>	Sep 22, 2015
<u>AB</u>		<u>10MG</u>	<u>A207416 002</u>	Sep 22, 2015
<u>AB</u>		<u>20MG</u>	<u>A207416 003</u>	Sep 22, 2015
<u>AB</u>	BIONPHARMA	<u>5MG</u>	<u>A209753 001</u>	Mar 02, 2018
<u>AB</u>		<u>10MG</u>	<u>A209753 002</u>	Mar 02, 2018
<u>AB</u>		<u>20MG</u>	<u>A209753 003</u>	Mar 02, 2018
<u>AB</u>	CEDIPROF INC	<u>5MG</u>	<u>A208737 001</u>	Feb 01, 2019
<u>AB</u>		<u>10MG</u>	<u>A208737 002</u>	Feb 01, 2019
<u>AB</u>		<u>20MG</u>	<u>A208737 003</u>	Feb 01, 2019
<u>AB</u>	MOUNTAIN	<u>5MG</u>	<u>A091159 001</u>	Mar 12, 2014
<u>AB</u>		<u>10MG</u>	<u>A091159 002</u>	Mar 12, 2014
<u>AB</u>		<u>20MG</u>	<u>A091159 003</u>	Mar 12, 2014
<u>AB</u>	NOVEL LABS INC	<u>5MG</u>	<u>A207884 001</u>	Nov 13, 2015
<u>AB</u>		<u>10MG</u>	<u>A207884 002</u>	Nov 13, 2015
<u>AB</u>		<u>20MG</u>	<u>A207884 003</u>	Nov 13, 2015
<u>AB</u>	OXFORD PHARMS	<u>5MG</u>	<u>A202892 001</u>	Sep 23, 2014
<u>AB</u>		<u>10MG</u>	<u>A202892 002</u>	Sep 23, 2014
<u>AB</u>		<u>20MG</u>	<u>A202892 003</u>	Sep 23, 2014
<u>AB</u>	PRINSTON INC	<u>5MG</u>	<u>A212697 001</u>	Jul 23, 2020
<u>AB</u>		<u>10MG</u>	<u>A212697 002</u>	Jul 23, 2020
<u>AB</u>		<u>20MG</u>	<u>A212697 003</u>	Jul 23, 2020
<u>AB</u>	SPECGX LLC	<u>5MG</u>	<u>A040300 001</u>	Nov 27, 1998
<u>AB</u>		<u>10MG</u>	<u>A040300 002</u>	Nov 27, 1998
<u>AB</u>		<u>20MG</u>	<u>A040300 003</u>	Nov 27, 1998
<u>AB</u>	SUN PHARM INDS INC	<u>5MG</u>	<u>A090710 001</u>	Mar 15, 2012
<u>AB</u>		<u>10MG</u>	<u>A090710 002</u>	Mar 15, 2012
<u>AB</u>		<u>20MG</u>	<u>A090710 003</u>	Mar 15, 2012

RITALIN

<u>AB</u>	+	SANDOZ	<u>5MG</u>	<u>N010187 003</u>
<u>AB</u>	+		<u>10MG</u>	<u>N010187 006</u>
<u>AB</u>	+	!	<u>20MG</u>	<u>N010187 010</u>

TABLET, CHEWABLE;ORAL

METHYLPHENIDATE HYDROCHLORIDE

<u>AB</u>	ASCENT PHARMS INC	<u>2.5MG</u>	<u>A210354 001</u>	Dec 29, 2017
<u>AB</u>		<u>5MG</u>	<u>A210354 002</u>	Dec 29, 2017
<u>AB</u>	!	<u>10MG</u>	<u>A210354 003</u>	Dec 29, 2017
<u>AB</u>	RISING	<u>2.5MG</u>	<u>A205756 001</u>	Nov 07, 2016
<u>AB</u>		<u>5MG</u>	<u>A205756 002</u>	Nov 07, 2016
<u>AB</u>		<u>10MG</u>	<u>A205756 003</u>	Nov 07, 2016

TABLET, EXTENDED RELEASE;ORAL

CONCERTA

<u>AB</u>	+	JANSSEN PHARMS	<u>18MG</u>	<u>N021121 001</u>	Aug 01, 2000
<u>AB</u>	+		<u>27MG</u>	<u>N021121 004</u>	Apr 01, 2002
<u>AB</u>	+		<u>36MG</u>	<u>N021121 002</u>	Aug 01, 2000
<u>AB</u>	+	!	<u>54MG</u>	<u>N021121 003</u>	Dec 08, 2000

METHYLIN ER

<u>AB</u>	SPECGX LLC	<u>10MG</u>	<u>A075629 001</u>	May 09, 2000
<u>AB</u>		<u>20MG</u>	<u>A075629 002</u>	May 09, 2000

METHYLPHENIDATE HYDROCHLORIDE

<u>AB</u>	ABHAI LLC	<u>10MG</u>	<u>A207488 001</u>	Jun 09, 2015
<u>AB</u>	!	<u>20MG</u>	<u>A207488 002</u>	Jun 09, 2015
<u>AB</u>	ACTAVIS LABS FL	<u>18MG</u>	<u>A076772 001</u>	Mar 22, 2018
<u>AB</u>		<u>27MG</u>	<u>A076772 002</u>	Mar 22, 2018
<u>AB</u>		<u>36MG</u>	<u>A076772 003</u>	Mar 22, 2018
<u>AB</u>		<u>54MG</u>	<u>A076655 001</u>	Mar 21, 2018

PRESCRIPTION DRUG PRODUCT LIST

METHYLPHENIDATE HYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL

METHYLPHENIDATE HYDROCHLORIDE

AB		72MG	A076655 002	Feb 28, 2022
AB	ALKEM LABS LTD	18MG	A214447 001	May 23, 2023
AB		20MG	A212288 001	Oct 06, 2020
AB		27MG	A214447 002	May 23, 2023
AB		36MG	A214447 003	May 23, 2023
AB		54MG	A214447 004	May 23, 2023
AB	ANDOR PHARMS	18MG	A211918 001	Apr 24, 2019
AB		27MG	A211918 002	Apr 24, 2019
AB		36MG	A211918 003	Apr 24, 2019
AB		54MG	A211918 004	Apr 24, 2019
AB	ASCENT PHARMS INC	18MG	A211009 001	Sep 03, 2019
AB		27MG	A211009 002	Sep 03, 2019
AB		36MG	A211009 003	Sep 03, 2019
AB		54MG	A211009 004	Sep 03, 2019
AB	AUROLIFE PHARMA LLC	18MG	A206726 001	Oct 21, 2016
AB		27MG	A206726 002	Oct 21, 2016
AB		36MG	A206726 003	Oct 21, 2016
AB		54MG	A206726 004	Oct 21, 2016
AB	DR REDDYS	18MG	A213473 001	Jul 29, 2020
AB		27MG	A213473 002	Jul 29, 2020
AB		36MG	A213473 003	Jul 29, 2020
AB		54MG	A213473 004	Jul 29, 2020
AB	GRANULES	10MG	A210992 001	Nov 21, 2018
AB		20MG	A210992 002	Nov 21, 2018
AB	OSMOTICA PHARM US	18MG	A205327 001	Jul 28, 2017
AB		27MG	A205327 002	Jul 28, 2017
AB		36MG	A205327 003	Jul 28, 2017
AB		54MG	A205327 004	Jul 28, 2017
AB	!	72MG	A205327 005	Jul 28, 2017
AB	SUN PHARM INDS INC	18MG	A205135 001	Aug 19, 2020
AB		27MG	A205135 002	Aug 19, 2020
AB		36MG	A205135 003	Aug 19, 2020
AB		54MG	A205135 004	Aug 19, 2020
AB		72MG	A217229 001	Aug 25, 2023
BX	LANNETT CO INC	18MG	A091695 001	Jul 09, 2013
BX		27MG	A091695 002	Jul 09, 2013
BX		36MG	A091695 003	Sep 23, 2013
BX		54MG	A091695 004	Sep 23, 2013
BX	SPECGX LLC	27MG	A202608 001	Dec 28, 2012
BX		36MG	A202608 002	Dec 28, 2012
BX		54MG	A202608 003	Dec 28, 2012
	RELEXXII			
+	OSMOTICA PHARM US	18MG	N216117 001	Jun 23, 2022
+		27MG	N216117 002	Jun 23, 2022
+		36MG	N216117 003	Jun 23, 2022
+		45MG	N216117 004	Jun 23, 2022
+		54MG	N216117 005	Jun 23, 2022
+		63MG	N216117 006	Jun 23, 2022
+	!	72MG	N216117 007	Jun 23, 2022

TABLET, EXTENDED RELEASE, CHEWABLE;ORAL

QUILLICHEW ER

+	NEXTWAVE PHARMS	20MG	N207960 001	Dec 04, 2015
+		30MG	N207960 002	Dec 04, 2015
+	!	40MG	N207960 003	Dec 04, 2015

METHYLPREDNISOLONE

TABLET;ORAL

MEDROL

AB	+	PFIZER	4MG	N011153 001
AB	+		8MG	N011153 004
AB	+		16MG	N011153 003
AB	+	!	32MG	N011153 006

METHYLPREDNISOLONE

AB	ENDO OPERATIONS	4MG	A040183 001	Dec 22, 1998
AB	JUBILANT CADISTA	4MG	A040189 001	Oct 31, 1997
AB		8MG	A040189 002	Oct 31, 1997
AB		16MG	A040189 003	Jul 20, 2007
AB		32MG	A040189 004	Jul 20, 2007
AB	SANDOZ	4MG	A040194 001	Oct 31, 1997
AB	TIANJIN TIANYAO	4MG	A204072 001	May 14, 2018

PRESCRIPTION DRUG PRODUCT LIST

METHYLPREDNISOLONE

TABLET; ORAL

METHYLPREDNISOLONE

<u>AB</u>	WATSON LABS	<u>4MG</u>	<u>A040232</u>	<u>001</u>	Oct 16, 1997
<u>AB</u>	ZYDUS PHARMS	<u>4MG</u>	<u>A206751</u>	<u>001</u>	Apr 23, 2018
<u>AB</u>		<u>8MG</u>	<u>A206751</u>	<u>002</u>	Apr 23, 2018
<u>AB</u>		<u>16MG</u>	<u>A206751</u>	<u>003</u>	Apr 23, 2018
<u>AB</u>		<u>32MG</u>	<u>A206751</u>	<u>004</u>	Apr 23, 2018
	MEDROL				
	+ PFIZER	2MG	N011153	002	

METHYLPREDNISOLONE ACETATE

INJECTABLE; INJECTION

DEPO-MEDROL

<u>AB</u>	+ PFIZER	<u>40MG/ML</u>	<u>N011757</u>	<u>001</u>	
<u>AB</u>	+	<u>80MG/ML</u>	<u>N011757</u>	<u>004</u>	

METHYLPREDNISOLONE ACETATE

<u>AB</u>	! AMNEAL	<u>40MG/ML</u>	<u>A210043</u>	<u>001</u>	May 20, 2019
<u>AB</u>		<u>40MG/ML</u>	<u>A216502</u>	<u>001</u>	Nov 03, 2023
<u>AB</u>	!	<u>80MG/ML</u>	<u>A210043</u>	<u>002</u>	May 20, 2019
<u>AB</u>		<u>80MG/ML</u>	<u>A216502</u>	<u>002</u>	Nov 03, 2023
<u>AB</u>	ENDO OPERATIONS	<u>40MG/ML</u>	<u>A214297</u>	<u>001</u>	Jan 21, 2022
<u>AB</u>		<u>80MG/ML</u>	<u>A214297</u>	<u>002</u>	Jan 21, 2022
<u>AB</u>	EUGIA PHARMA	<u>40MG/ML</u>	<u>A211930</u>	<u>001</u>	Apr 13, 2023
<u>AB</u>		<u>80MG/ML</u>	<u>A211930</u>	<u>002</u>	Apr 13, 2023
<u>AB</u>	HONG KONG	<u>40MG/ML</u>	<u>A040557</u>	<u>001</u>	Feb 23, 2005
<u>AB</u>		<u>80MG/ML</u>	<u>A040557</u>	<u>002</u>	Feb 23, 2005
<u>AB</u>	! SAGENT PHARMS INC	<u>40MG/ML</u>	<u>A201835</u>	<u>002</u>	Jun 27, 2018
<u>AB</u>	!	<u>80MG/ML</u>	<u>A201835</u>	<u>003</u>	Jun 27, 2018
<u>AB</u>	SANDOZ	<u>40MG/ML</u>	<u>A040719</u>	<u>001</u>	Jan 29, 2009
<u>AB</u>		<u>40MG/ML</u>	<u>A040794</u>	<u>001</u>	Mar 05, 2009
<u>AB</u>		<u>80MG/ML</u>	<u>A040719</u>	<u>002</u>	Jan 29, 2009
<u>AB</u>		<u>80MG/ML</u>	<u>A040794</u>	<u>002</u>	Mar 05, 2009
<u>AB</u>	WILSHIRE PHARMS INC	<u>40MG/ML</u>	<u>A214870</u>	<u>002</u>	May 10, 2023
<u>AB</u>		<u>80MG/ML</u>	<u>A214870</u>	<u>001</u>	Jan 03, 2023
	DEPO-MEDROL				
	+! PFIZER	20MG/ML	N011757	002	

METHYLPREDNISOLONE SODIUM SUCCINATE

INJECTABLE; INJECTION

METHYLPREDNISOLONE SODIUM SUCCINATE

<u>AP</u>	AMNEAL	<u>EQ 40MG BASE/VIAL</u>	<u>A207549</u>	<u>001</u>	Nov 09, 2016
<u>AP</u>		<u>EQ 125MG BASE/VIAL</u>	<u>A207549</u>	<u>002</u>	Nov 09, 2016
<u>AP</u>	FRESENIUS KABI USA	<u>EQ 40MG BASE/VIAL</u>	<u>A040583</u>	<u>001</u>	Jul 30, 2004
<u>AP</u>		<u>EQ 125MG BASE/VIAL</u>	<u>A040583</u>	<u>002</u>	Jul 30, 2004
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>A040612</u>	<u>001</u>	Aug 12, 2004
<u>AP</u>	HIKMA	<u>EQ 40MG BASE/VIAL</u>	<u>A203125</u>	<u>001</u>	Sep 26, 2022
<u>AP</u>		<u>EQ 125MG BASE/VIAL</u>	<u>A203125</u>	<u>002</u>	Sep 26, 2022
<u>AP</u>		<u>EQ 500MG BASE/VIAL</u>	<u>A202691</u>	<u>001</u>	Feb 16, 2016
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>A202691</u>	<u>002</u>	Feb 16, 2016
<u>AP</u>	SAGENT PHARMS INC	<u>EQ 40MG BASE/VIAL</u>	<u>A040888</u>	<u>001</u>	Jul 18, 2011
<u>AP</u>		<u>EQ 125MG BASE/VIAL</u>	<u>A040888</u>	<u>002</u>	Jul 18, 2011
<u>AP</u>		<u>EQ 500MG BASE/VIAL</u>	<u>A040888</u>	<u>003</u>	Jul 18, 2011
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>A040888</u>	<u>004</u>	Jul 18, 2011
<u>AP</u>		<u>EQ 2GM BASE/VIAL</u>	<u>A040888</u>	<u>005</u>	Jul 18, 2011
<u>AP</u>	TIANJIN KINGYORK	<u>EQ 40MG BASE/VIAL</u>	<u>A212396</u>	<u>001</u>	Apr 20, 2021
<u>AP</u>		<u>EQ 125MG BASE/VIAL</u>	<u>A212396</u>	<u>002</u>	Apr 20, 2021
<u>AP</u>		<u>EQ 500MG BASE/VIAL</u>	<u>A212396</u>	<u>003</u>	Apr 20, 2021
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>A212396</u>	<u>004</u>	Apr 20, 2021
<u>AP</u>		<u>EQ 2GM BASE/VIAL</u>	<u>A212396</u>	<u>005</u>	Apr 20, 2021
	<u>SOLU-MEDROL</u>				
<u>AP</u>	+! PHARMACIA AND UPJOHN	<u>EQ 40MG BASE/VIAL</u>	<u>N011856</u>	<u>003</u>	
<u>AP</u>	+!	<u>EQ 125MG BASE/VIAL</u>	<u>N011856</u>	<u>004</u>	
<u>AP</u>	+!	<u>EQ 500MG BASE/VIAL</u>	<u>N011856</u>	<u>005</u>	
<u>AP</u>	+!	<u>EQ 1GM BASE/VIAL</u>	<u>N011856</u>	<u>006</u>	
<u>AP</u>	+!	<u>EQ 2GM BASE/VIAL</u>	<u>N011856</u>	<u>007</u>	Feb 27, 1985

PRESCRIPTION DRUG PRODUCT LIST

METHYLTESTOSTERONE

CAPSULE; ORAL

METHYLTESTOSTERONE

AB	!	IMPAX LABS INC	10MG	A204851	001	Sep 21, 2015
AB		NOVITIUM PHARMA	10MG	A215270	001	Feb 18, 2022

TABLET; ORAL

ANDROID 25

BP		VALEANT PHARM INTL	25MG	A087147	001	
----	--	--------------------	------	---------	-----	--

METHYLTESTOSTERONE

BP		IMPAX LABS	10MG	A080767	002	
----	--	------------	------	---------	-----	--

METOCLOPRAMIDE HYDROCHLORIDE

INJECTABLE; INJECTION

METOCLOPRAMIDE

AP		AVET LIFESCIENCES	EQ 5MG BASE/ML	A204756	001	Dec 20, 2013
-----------	--	-------------------	-----------------------	----------------	------------	--------------

METOCLOPRAMIDE HYDROCHLORIDE

AP		FRESENIUS KABI USA	EQ 5MG BASE/ML	A091392	001	Apr 19, 2013
-----------	--	--------------------	-----------------------	----------------	------------	--------------

AP	!	HOSPIRA	EQ 5MG BASE/ML	A073118	001	Jan 17, 1991
-----------	----------	---------	-----------------------	----------------	------------	--------------

AP		TEVA PHARMS USA	EQ 5MG BASE/ML	A073135	001	Nov 27, 1991
-----------	--	-----------------	-----------------------	----------------	------------	--------------

SOLUTION; ORAL

METOCLOPRAMIDE HYDROCHLORIDE

AA		ANI PHARMS	EQ 5MG BASE/5ML	A071402	001	Jun 25, 1993
-----------	--	------------	------------------------	----------------	------------	--------------

AA	!	GENUS	EQ 5MG BASE/5ML	A072744	001	May 28, 1991
-----------	----------	-------	------------------------	----------------	------------	--------------

SPRAY, METERED; NASAL

GIMOTI

+! EVOKE PHARMA INC

EQ 15MG BASE/SPRAY

N209388 001 Jun 19, 2020

TABLET; ORAL

METOCLOPRAMIDE HYDROCHLORIDE

AB		AIPING PHARM INC	EQ 5MG BASE	A072215	002	Nov 05, 2024
-----------	--	------------------	--------------------	----------------	------------	--------------

AB			EQ 10MG BASE	A072215	001	Jan 30, 1990
-----------	--	--	---------------------	----------------	------------	--------------

AB		IMPAX LABS INC	EQ 5MG BASE	A071250	002	Dec 28, 1995
-----------	--	----------------	--------------------	----------------	------------	--------------

AB			EQ 10MG BASE	A071250	001	Feb 03, 1988
-----------	--	--	---------------------	----------------	------------	--------------

AB		IPCA LABS LTD	EQ 5MG BASE	A078807	001	Jun 12, 2008
-----------	--	---------------	--------------------	----------------	------------	--------------

AB			EQ 10MG BASE	A078807	002	Jun 12, 2008
-----------	--	--	---------------------	----------------	------------	--------------

AB		STRIDES PHARMA	EQ 5MG BASE	A077878	001	Aug 28, 2006
-----------	--	----------------	--------------------	----------------	------------	--------------

AB			EQ 10MG BASE	A070581	001	Oct 17, 1985
-----------	--	--	---------------------	----------------	------------	--------------

AB			EQ 10MG BASE	A077878	002	Aug 28, 2006
-----------	--	--	---------------------	----------------	------------	--------------

AB		TEVA	EQ 5MG BASE	A072801	001	Jun 15, 1993
-----------	--	------	--------------------	----------------	------------	--------------

AB			EQ 10MG BASE	A070184	001	Jul 29, 1985
-----------	--	--	---------------------	----------------	------------	--------------

REGLAN

AB	+	ANI PHARMS	EQ 5MG BASE	N017854	002	May 05, 1987
-----------	----------	------------	--------------------	----------------	------------	--------------

AB	+	!	EQ 10MG BASE	N017854	001	
-----------	----------	----------	---------------------	----------------	------------	--

TABLET, ORALLY DISINTEGRATING; ORAL

METOCLOPRAMIDE HYDROCHLORIDE

NOVEL LABS INC

EQ 5MG BASE

A202191 001 Aug 15, 2014

!

EQ 10MG BASE

A202191 002 Aug 15, 2014

METOLAZONE

TABLET; ORAL

METOLAZONE

AB		ALEMBIC	2.5MG	A213251	001	Dec 02, 2020
-----------	--	---------	--------------	----------------	------------	--------------

AB			5MG	A213251	002	Dec 02, 2020
-----------	--	--	------------	----------------	------------	--------------

AB			10MG	A213251	003	Dec 02, 2020
-----------	--	--	-------------	----------------	------------	--------------

AB		CHEMISTRY HLTH	2.5MG	A218606	001	Apr 17, 2024
-----------	--	----------------	--------------	----------------	------------	--------------

AB			5MG	A218606	002	Apr 17, 2024
-----------	--	--	------------	----------------	------------	--------------

AB			10MG	A218606	003	Apr 17, 2024
-----------	--	--	-------------	----------------	------------	--------------

AB		INNOGENIX	2.5MG	A213827	001	Mar 30, 2021
-----------	--	-----------	--------------	----------------	------------	--------------

AB			5MG	A213827	002	Mar 30, 2021
-----------	--	--	------------	----------------	------------	--------------

AB			10MG	A213827	003	Mar 30, 2021
-----------	--	--	-------------	----------------	------------	--------------

AB		MICRO LABS	2.5MG	A217563	001	Nov 06, 2024
-----------	--	------------	--------------	----------------	------------	--------------

AB			5MG	A217563	002	Nov 06, 2024
-----------	--	--	------------	----------------	------------	--------------

AB			10MG	A217563	003	Nov 06, 2024
-----------	--	--	-------------	----------------	------------	--------------

AB		MYLAN	2.5MG	A076698	001	Dec 23, 2003
-----------	--	-------	--------------	----------------	------------	--------------

AB			5MG	A076698	002	Oct 19, 2004
-----------	--	--	------------	----------------	------------	--------------

AB			10MG	A076698	003	Oct 19, 2004
-----------	--	--	-------------	----------------	------------	--------------

AB		NE RX PHARMA	2.5MG	A216216	001	Oct 25, 2022
-----------	--	--------------	--------------	----------------	------------	--------------

AB			5MG	A216216	002	Oct 25, 2022
-----------	--	--	------------	----------------	------------	--------------

AB			10MG	A216216	003	Oct 25, 2022
-----------	--	--	-------------	----------------	------------	--------------

AB		RENATA	2.5MG	A215616	001	Oct 24, 2022
-----------	--	--------	--------------	----------------	------------	--------------

AB			5MG	A215616	002	Oct 24, 2022
-----------	--	--	------------	----------------	------------	--------------

AB			10MG	A215616	003	Oct 24, 2022
-----------	--	--	-------------	----------------	------------	--------------

AB		RUBICON	2.5MG	A215184	001	Aug 20, 2021
-----------	--	---------	--------------	----------------	------------	--------------

AB			5MG	A215184	002	Aug 20, 2021
-----------	--	--	------------	----------------	------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

METOLAZONE

TABLET;ORAL

METOLAZONE

<u>AB</u>		<u>10MG</u>	<u>A215184</u>	<u>003</u>	Aug 20, 2021
<u>AB</u>	SANDOZ	<u>2.5MG</u>	<u>A076732</u>	<u>001</u>	Dec 19, 2003
<u>AB</u>	!	<u>5MG</u>	<u>A076466</u>	<u>001</u>	Dec 19, 2003
<u>AB</u>	!	<u>10MG</u>	<u>A076466</u>	<u>002</u>	Dec 19, 2003
<u>AB</u>	UNICHEM	<u>2.5MG</u>	<u>A214799</u>	<u>001</u>	Mar 30, 2021
<u>AB</u>		<u>5MG</u>	<u>A214799</u>	<u>002</u>	Mar 30, 2021
<u>AB</u>		<u>10MG</u>	<u>A214799</u>	<u>003</u>	Mar 30, 2021

METOPROLOL SUCCINATE

CAPSULE, EXTENDED RELEASE;ORAL

KAPSPARGO SPRINKLE

+	SPIL	EQ 25MG TARTRATE	N210428	001	Jan 26, 2018
+		EQ 50MG TARTRATE	N210428	002	Jan 26, 2018
+		EQ 100MG TARTRATE	N210428	003	Jan 26, 2018
+	!	EQ 200MG TARTRATE	N210428	004	Jan 26, 2018

TABLET, EXTENDED RELEASE;ORAL

METOPROLOL SUCCINATE

<u>AB</u>	ACTAVIS ELIZABETH	<u>EQ 25MG TARTRATE</u>	<u>A204161</u>	<u>001</u>	Nov 25, 2016
<u>AB</u>		<u>EQ 50MG TARTRATE</u>	<u>A204161</u>	<u>002</u>	Nov 25, 2016
<u>AB</u>		<u>EQ 100MG TARTRATE</u>	<u>A204161</u>	<u>003</u>	Nov 25, 2016
<u>AB</u>		<u>EQ 200MG TARTRATE</u>	<u>A204161</u>	<u>004</u>	Nov 25, 2016
<u>AB</u>	ACTAVIS LABS FL INC	<u>EQ 50MG TARTRATE</u>	<u>A076862</u>	<u>001</u>	Aug 03, 2009
<u>AB</u>	ALKEM LABS LTD	<u>EQ 25MG TARTRATE</u>	<u>A211143</u>	<u>001</u>	Nov 25, 2020
<u>AB</u>		<u>EQ 50MG TARTRATE</u>	<u>A211143</u>	<u>002</u>	Nov 25, 2020
<u>AB</u>		<u>EQ 100MG TARTRATE</u>	<u>A211143</u>	<u>003</u>	Nov 25, 2020
<u>AB</u>		<u>EQ 200MG TARTRATE</u>	<u>A211143</u>	<u>004</u>	Nov 25, 2020
<u>AB</u>	CIPLA	<u>EQ 50MG TARTRATE</u>	<u>A207465</u>	<u>001</u>	Oct 26, 2018
<u>AB</u>		<u>EQ 100MG TARTRATE</u>	<u>A207465</u>	<u>002</u>	Oct 26, 2018
<u>AB</u>		<u>EQ 200MG TARTRATE</u>	<u>A207465</u>	<u>003</u>	Oct 26, 2018
<u>AB</u>	DR REDDYS LABS LTD	<u>EQ 25MG TARTRATE</u>	<u>A090617</u>	<u>001</u>	Aug 01, 2012
<u>AB</u>		<u>EQ 50MG TARTRATE</u>	<u>A090617</u>	<u>002</u>	Aug 01, 2012
<u>AB</u>	GRANULES	<u>EQ 25MG TARTRATE</u>	<u>A216916</u>	<u>001</u>	Jun 12, 2023
<u>AB</u>		<u>EQ 50MG TARTRATE</u>	<u>A216916</u>	<u>002</u>	Jun 12, 2023
<u>AB</u>		<u>EQ 100MG TARTRATE</u>	<u>A216916</u>	<u>003</u>	Jun 12, 2023
<u>AB</u>		<u>EQ 200MG TARTRATE</u>	<u>A216916</u>	<u>004</u>	Jun 12, 2023
<u>AB</u>	HETERO LABS LTD III	<u>EQ 25MG TARTRATE</u>	<u>A205541</u>	<u>001</u>	Nov 06, 2020
<u>AB</u>		<u>EQ 50MG TARTRATE</u>	<u>A205541</u>	<u>002</u>	Nov 06, 2020
<u>AB</u>		<u>EQ 100MG TARTRATE</u>	<u>A205541</u>	<u>003</u>	Nov 06, 2020
<u>AB</u>		<u>EQ 200MG TARTRATE</u>	<u>A205541</u>	<u>004</u>	Nov 06, 2020
<u>AB</u>	MYLAN PHARMS INC	<u>EQ 25MG TARTRATE</u>	<u>A202033</u>	<u>001</u>	Dec 15, 2011
<u>AB</u>		<u>EQ 50MG TARTRATE</u>	<u>A202033</u>	<u>002</u>	Dec 15, 2011
<u>AB</u>		<u>EQ 100MG TARTRATE</u>	<u>A202033</u>	<u>003</u>	Dec 15, 2011
<u>AB</u>		<u>EQ 200MG TARTRATE</u>	<u>A202033</u>	<u>004</u>	Dec 15, 2011
<u>AB</u>	NOVAST LABS	<u>EQ 25MG TARTRATE</u>	<u>A204106</u>	<u>001</u>	Feb 06, 2018
<u>AB</u>		<u>EQ 50MG TARTRATE</u>	<u>A204106</u>	<u>002</u>	Feb 06, 2018
<u>AB</u>		<u>EQ 100MG TARTRATE</u>	<u>A204106</u>	<u>003</u>	Feb 06, 2018
<u>AB</u>		<u>EQ 200MG TARTRATE</u>	<u>A204106</u>	<u>004</u>	Feb 06, 2018
<u>AB</u>	PHARMADAX INC	<u>EQ 25MG TARTRATE</u>	<u>A203028</u>	<u>001</u>	Mar 31, 2020
<u>AB</u>		<u>EQ 50MG TARTRATE</u>	<u>A203028</u>	<u>002</u>	Mar 31, 2020
<u>AB</u>		<u>EQ 100MG TARTRATE</u>	<u>A203699</u>	<u>001</u>	Mar 30, 2020
<u>AB</u>		<u>EQ 200MG TARTRATE</u>	<u>A203699</u>	<u>002</u>	Mar 30, 2020
<u>AB</u>	REDDYS	<u>EQ 100MG TARTRATE</u>	<u>A078889</u>	<u>001</u>	Aug 15, 2012
<u>AB</u>		<u>EQ 200MG TARTRATE</u>	<u>A078889</u>	<u>002</u>	Aug 15, 2012
<u>AB</u>	SUNSHINE	<u>EQ 25MG TARTRATE</u>	<u>A214004</u>	<u>004</u>	May 23, 2024
<u>AB</u>		<u>EQ 50MG TARTRATE</u>	<u>A214004</u>	<u>001</u>	Sep 26, 2022
<u>AB</u>		<u>EQ 100MG TARTRATE</u>	<u>A214004</u>	<u>002</u>	Sep 26, 2022
<u>AB</u>		<u>EQ 200MG TARTRATE</u>	<u>A214004</u>	<u>003</u>	Sep 26, 2022
<u>AB</u>	VISUM PHARM	<u>EQ 25MG TARTRATE</u>	<u>A207206</u>	<u>001</u>	Dec 19, 2018
<u>AB</u>		<u>EQ 50MG TARTRATE</u>	<u>A207206</u>	<u>002</u>	Dec 19, 2018
<u>AB</u>		<u>EQ 100MG TARTRATE</u>	<u>A207206</u>	<u>003</u>	Dec 19, 2018
<u>AB</u>		<u>EQ 200MG TARTRATE</u>	<u>A207206</u>	<u>004</u>	Dec 19, 2018
<u>AB</u>	WOCKHARDT	<u>EQ 25MG TARTRATE</u>	<u>A090615</u>	<u>001</u>	Jul 22, 2010
<u>AB</u>		<u>EQ 50MG TARTRATE</u>	<u>A090615</u>	<u>002</u>	Jul 22, 2010
<u>AB</u>		<u>EQ 100MG TARTRATE</u>	<u>A090615</u>	<u>003</u>	Jul 22, 2010
<u>AB</u>		<u>EQ 200MG TARTRATE</u>	<u>A090615</u>	<u>004</u>	Jul 22, 2010
<u>AB</u>	YICHANG HUMANWELL	<u>EQ 25MG TARTRATE</u>	<u>A213854</u>	<u>004</u>	Aug 01, 2022
<u>AB</u>		<u>EQ 50MG TARTRATE</u>	<u>A213854</u>	<u>003</u>	Nov 08, 2021
<u>AB</u>		<u>EQ 100MG TARTRATE</u>	<u>A213854</u>	<u>001</u>	Feb 12, 2021
<u>AB</u>		<u>EQ 200MG TARTRATE</u>	<u>A213854</u>	<u>002</u>	Feb 12, 2021
<u>AB</u>	ZHEJIANG JUTAI	<u>EQ 50MG TARTRATE</u>	<u>A214110</u>	<u>001</u>	Jul 26, 2024

PRESCRIPTION DRUG PRODUCT LIST

METOPROLOL SUCCINATE

TABLET, EXTENDED RELEASE;ORAL

METOPROLOL SUCCINATE

PHARM

<u>AB</u>		<u>EQ 100MG TARTRATE</u>	<u>A214110 002</u>	Jul 26, 2024
<u>AB</u>		<u>EQ 200MG TARTRATE</u>	<u>A214110 003</u>	Jul 26, 2024
<u>AB</u>	ZYDUS PHARMS	<u>EQ 25MG TARTRATE</u>	<u>A203894 001</u>	Mar 23, 2018
<u>AB</u>		<u>EQ 50MG TARTRATE</u>	<u>A203894 002</u>	Mar 23, 2018
<u>AB</u>		<u>EQ 100MG TARTRATE</u>	<u>A203894 003</u>	Mar 23, 2018
<u>AB</u>		<u>EQ 200MG TARTRATE</u>	<u>A203894 004</u>	Mar 23, 2018

TOPROL-XL

<u>AB</u>	+	TOPROL	<u>EQ 25MG TARTRATE</u>	<u>N019962 004</u>	Feb 05, 2001
<u>AB</u>	+	!	<u>EQ 50MG TARTRATE</u>	<u>N019962 001</u>	Jan 10, 1992
<u>AB</u>	+		<u>EQ 100MG TARTRATE</u>	<u>N019962 002</u>	Jan 10, 1992
<u>AB</u>	+	!	<u>EQ 200MG TARTRATE</u>	<u>N019962 003</u>	Jan 10, 1992

METOPROLOL TARTRATE

INJECTABLE; INJECTION

METOPROLOL TARTRATE

<u>AP</u>		BAXTER HLTHCARE CORP	<u>1MG/ML</u>	<u>A078950 001</u>	Apr 29, 2013
<u>AP</u>		FRESENIUS KABI USA	<u>1MG/ML</u>	<u>A091045 001</u>	Oct 25, 2010
<u>AP</u>		GLAND PHARMA LTD	<u>1MG/ML</u>	<u>A204205 001</u>	Aug 27, 2014
<u>AP</u>		HIKMA	<u>1MG/ML</u>	<u>A076495 001</u>	Jul 07, 2003
<u>AP</u>		HIKMA FARMACEUTICA	<u>1MG/ML</u>	<u>A077761 001</u>	May 30, 2007
<u>AP</u>	!	HOSPIRA	<u>1MG/ML</u>	<u>A078085 001</u>	Apr 29, 2008
<u>AP</u>		SANDOZ	<u>1MG/ML</u>	<u>A077360 001</u>	Oct 02, 2007

TABLET;ORAL

LOPRESSOR

<u>AB</u>	+	VALIDUS PHARMS	<u>50MG</u>	<u>N017963 001</u>	
<u>AB</u>	+		<u>100MG</u>	<u>N017963 002</u>	

METOPROLOL TARTRATE

<u>AB</u>		ALEMBIC PHARMS LTD	<u>25MG</u>	<u>A202871 001</u>	May 28, 2013
<u>AB</u>			<u>50MG</u>	<u>A202871 002</u>	May 28, 2013
<u>AB</u>			<u>100MG</u>	<u>A202871 003</u>	May 28, 2013
<u>AB</u>		AUROBINDO PHARMA	<u>25MG</u>	<u>A077739 001</u>	Sep 11, 2007
<u>AB</u>			<u>37.5MG</u>	<u>A077739 004</u>	Dec 07, 2023
<u>AB</u>			<u>50MG</u>	<u>A077739 002</u>	Sep 11, 2007
<u>AB</u>			<u>75MG</u>	<u>A077739 005</u>	Dec 07, 2023
<u>AB</u>			<u>100MG</u>	<u>A077739 003</u>	Sep 11, 2007
<u>AB</u>		IPCA LABS LTD	<u>25MG</u>	<u>A078459 001</u>	Jun 17, 2008
<u>AB</u>			<u>50MG</u>	<u>A078459 002</u>	Jun 17, 2008
<u>AB</u>			<u>100MG</u>	<u>A078459 003</u>	Jun 17, 2008
<u>AB</u>		MYLAN	<u>25MG</u>	<u>A076704 001</u>	Jan 16, 2004
<u>AB</u>			<u>37.5MG</u>	<u>A076704 004</u>	Mar 18, 2015
<u>AB</u>			<u>50MG</u>	<u>A076704 002</u>	Jan 16, 2004
<u>AB</u>			<u>75MG</u>	<u>A076704 005</u>	Mar 18, 2015
<u>AB</u>	!		<u>100MG</u>	<u>A076704 003</u>	Jan 16, 2004
<u>AB</u>		RENATA	<u>50MG</u>	<u>A074453 001</u>	Apr 27, 1995
<u>AB</u>			<u>100MG</u>	<u>A074453 002</u>	Apr 27, 1995
<u>AB</u>		RUBICON	<u>25MG</u>	<u>A200981 001</u>	Oct 28, 2014
<u>AB</u>			<u>37.5MG</u>	<u>A200981 004</u>	Aug 21, 2019
<u>AB</u>			<u>50MG</u>	<u>A200981 002</u>	Oct 28, 2014
<u>AB</u>			<u>75MG</u>	<u>A200981 005</u>	Aug 21, 2019
<u>AB</u>			<u>100MG</u>	<u>A200981 003</u>	Oct 28, 2014
<u>AB</u>		SUN PHARM INDS INC	<u>25MG</u>	<u>A076670 001</u>	Jan 15, 2004
<u>AB</u>			<u>50MG</u>	<u>A074644 001</u>	Dec 10, 1996
<u>AB</u>			<u>100MG</u>	<u>A074644 002</u>	Dec 10, 1996
<u>AB</u>		YOUNGTECH PHARMS INC	<u>25MG</u>	<u>A208955 001</u>	Feb 05, 2020
<u>AB</u>			<u>50MG</u>	<u>A208955 002</u>	Feb 05, 2020
<u>AB</u>			<u>100MG</u>	<u>A208955 003</u>	Feb 05, 2020

METRONIDAZOLE

CAPSULE;ORAL

FLAGYL

<u>AB</u>	+	!	PFIZER	<u>375MG</u>	<u>N020334 001</u>	May 03, 1995
-----------	---	---	--------	--------------	--------------------	--------------

METRONIDAZOLE

<u>AB</u>		ALEMBIC	<u>375MG</u>	<u>A079065 001</u>	Jun 23, 2009
<u>AB</u>		CHARTWELL RX	<u>375MG</u>	<u>A076522 001</u>	Jan 29, 2004

CREAM; TOPICAL

METROCREAM

<u>AB</u>	+	!	GALDERMA LABS LP	<u>0.75%</u>	<u>N020531 001</u>	Sep 20, 1995
-----------	---	---	------------------	--------------	--------------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

METRONIDAZOLE

CREAM; TOPICAL

METRONIDAZOLE

<u>AB</u>	COSETTE	<u>0.75%</u>	<u>A077549</u>	<u>001</u>	Dec 19, 2007
<u>AB</u>	FOUGERA PHARMS	<u>0.75%</u>	<u>A076408</u>	<u>001</u>	May 28, 2004
<u>AB</u>	ZYDUS LIFESCIENCES	<u>0.75%</u>	<u>A217128</u>	<u>001</u>	Apr 21, 2023

NORITATE

+	BAUSCH	1%	N020743	001	Sep 26, 1997
---	--------	----	---------	-----	--------------

GEL; TOPICAL

METROGEL

<u>AB</u>	+	GALDERMA LABS LP	<u>1%</u>	<u>N021789</u>	<u>001</u>	Jun 30, 2005
-----------	---	------------------	-----------	----------------	------------	--------------

METRONIDAZOLE

<u>AB</u>		ALEMBIC	<u>1%</u>	<u>A212646</u>	<u>001</u>	Sep 03, 2021
<u>AB</u>		AUROBINDO PHARMA LTD	<u>1%</u>	<u>A218941</u>	<u>001</u>	Oct 15, 2024
<u>AB</u>	!	COSETTE	<u>0.75%</u>	<u>A078178</u>	<u>001</u>	Jan 19, 2011
<u>AB</u>			<u>1%</u>	<u>A216692</u>	<u>001</u>	Jan 23, 2023
<u>AB</u>		ENCUBE	<u>0.75%</u>	<u>A077547</u>	<u>001</u>	Jul 13, 2006
<u>AB</u>		FOUGERA PHARMS	<u>0.75%</u>	<u>A077018</u>	<u>001</u>	Jun 06, 2006
<u>AB</u>		TARO	<u>0.75%</u>	<u>A077819</u>	<u>001</u>	Jul 18, 2006
<u>AB</u>			<u>1%</u>	<u>A204651</u>	<u>001</u>	Mar 14, 2017

GEL; VAGINAL

METROGEL-VAGINAL

<u>AB</u>	+	BAUSCH	<u>0.75%</u>	<u>N020208</u>	<u>001</u>	Aug 17, 1992
-----------	---	--------	--------------	----------------	------------	--------------

METRONIDAZOLE

<u>AB</u>		COSETTE	<u>0.75%</u>	<u>A216323</u>	<u>001</u>	Jul 12, 2023
<u>AB</u>		ENCUBE	<u>0.75%</u>	<u>A215610</u>	<u>001</u>	Jun 16, 2023
<u>AB</u>			<u>1.3%</u>	<u>A216795</u>	<u>001</u>	Mar 18, 2024
<u>AB</u>		GLENMARK PHARMS LTD	<u>0.75%</u>	<u>A215794</u>	<u>001</u>	Jan 27, 2022
<u>AB</u>		PADAGIS ISRAEL	<u>0.75%</u>	<u>A211786</u>	<u>001</u>	Jul 02, 2019
<u>AB</u>		SAPTALIS PHARMS	<u>0.75%</u>	<u>A216750</u>	<u>001</u>	Dec 31, 2024
<u>AB</u>		SOLARIS PHARMA CORP	<u>0.75%</u>	<u>A213648</u>	<u>001</u>	Oct 14, 2021

NUVESSA

<u>AB</u>	+	CHEMO RESEARCH SL	<u>1.3%</u>	<u>N205223</u>	<u>001</u>	Mar 24, 2014
-----------	---	-------------------	-------------	----------------	------------	--------------

VANDAZOLE

BX	!	TEVA PHARMS	0.75%	N021806	001	May 20, 2005
----	---	-------------	-------	---------	-----	--------------

INJECTABLE; INJECTION

FLAGYL I.V. RTU IN PLASTIC CONTAINER

<u>AP</u>	+	BAXTER HLTHCARE	<u>500MG/100ML</u>	<u>N018657</u>	<u>001</u>	
-----------	---	-----------------	--------------------	----------------	------------	--

METRO I.V. IN PLASTIC CONTAINER

<u>AP</u>	+	B BRAUN	<u>500MG/100ML</u>	<u>N018900</u>	<u>001</u>	Sep 29, 1983
-----------	---	---------	--------------------	----------------	------------	--------------

METRONIDAZOLE IN PLASTIC CONTAINER

<u>AP</u>		AMNEAL	<u>500MG/100ML</u>	<u>A217665</u>	<u>001</u>	May 24, 2023
<u>AP</u>		BAXTER HLTHCARE CORP	<u>500MG/100ML</u>	<u>A078084</u>	<u>001</u>	Mar 31, 2008
<u>AP</u>		GLAND PHARMA LTD	<u>500MG/100ML</u>	<u>A212435</u>	<u>001</u>	Aug 03, 2020
<u>AP</u>	+	HOSPIRA	<u>500MG/100ML</u>	<u>N018890</u>	<u>002</u>	Nov 18, 1983
<u>AP</u>		INFORLIFE	<u>500MG/100ML</u>	<u>A206191</u>	<u>001</u>	Feb 25, 2019

LOTION; TOPICAL

METROLOTION

<u>AB</u>	+	GALDERMA LABS LP	<u>0.75%</u>	<u>N020901</u>	<u>001</u>	Nov 24, 1998
-----------	---	------------------	--------------	----------------	------------	--------------

METRONIDAZOLE

<u>AB</u>		FOUGERA PHARMS	<u>0.75%</u>	<u>A077197</u>	<u>001</u>	May 24, 2006
-----------	--	----------------	--------------	----------------	------------	--------------

SUSPENSION; ORAL

LIKMEZ

+	SAPTALIS PHARMS	500MG/5ML	N216755	001	Sep 22, 2023
---	-----------------	-----------	---------	-----	--------------

TABLET; ORAL

METRONIDAZOLE

<u>AB</u>		ALEMBIC	<u>250MG</u>	<u>A208681</u>	<u>001</u>	Jun 20, 2017
<u>AB</u>			<u>500MG</u>	<u>A208681</u>	<u>002</u>	Jun 20, 2017
<u>AB</u>		ALEMBIC PHARMS LTD	<u>250MG</u>	<u>A079067</u>	<u>001</u>	Mar 13, 2009
<u>AB</u>			<u>500MG</u>	<u>A079067</u>	<u>002</u>	Mar 13, 2009
<u>AB</u>		APPCO	<u>250MG</u>	<u>A205245</u>	<u>001</u>	Sep 23, 2015
<u>AB</u>			<u>500MG</u>	<u>A205245</u>	<u>002</u>	Sep 23, 2015
<u>AB</u>		AUROBINDO PHARMA LTD	<u>250MG</u>	<u>A203974</u>	<u>001</u>	May 29, 2015
<u>AB</u>	!		<u>500MG</u>	<u>A203974</u>	<u>002</u>	May 29, 2015
<u>AB</u>		CADILA	<u>250MG</u>	<u>A206560</u>	<u>001</u>	Nov 16, 2016
<u>AB</u>			<u>500MG</u>	<u>A206560</u>	<u>002</u>	Nov 16, 2016
<u>AB</u>		CADILA PHARMS LTD	<u>250MG</u>	<u>A209794</u>	<u>001</u>	Dec 12, 2017
<u>AB</u>			<u>500MG</u>	<u>A209794</u>	<u>002</u>	Dec 12, 2017
<u>AB</u>		ESJAY PHARMA	<u>250MG</u>	<u>A208162</u>	<u>001</u>	May 25, 2016
<u>AB</u>			<u>500MG</u>	<u>A208162</u>	<u>002</u>	May 25, 2016

PRESCRIPTION DRUG PRODUCT LIST

METRONIDAZOLE

TABLET; ORAL

METRONIDAZOLE

<u>AB</u>	INNOGENIX	<u>250MG</u>	<u>A070772 001</u>	Jul 16, 1986
<u>AB</u>		<u>500MG</u>	<u>A070772 002</u>	Jul 16, 1986
<u>AB</u>	LUPIN LTD	<u>250MG</u>	<u>A209096 001</u>	Sep 12, 2017
<u>AB</u>		<u>500MG</u>	<u>A209096 002</u>	Sep 12, 2017
<u>AB</u>	PLIVA	<u>500MG</u>	<u>A070033 001</u>	Dec 06, 1984
<u>AB</u>	STRIDES PHARMA	<u>250MG</u>	<u>A070040 001</u>	Jan 29, 1985
<u>AB</u>		<u>500MG</u>	<u>A070039 001</u>	Jan 29, 1985
<u>AB</u>	TEVA PHARMS USA	<u>250MG</u>	<u>A070027 001</u>	Nov 06, 1984
<u>AB</u>	UNICHEM	<u>250MG</u>	<u>A203458 001</u>	Jan 22, 2014
<u>AB</u>		<u>500MG</u>	<u>A203458 002</u>	Jan 22, 2014
<u>AB</u>	WATSON LABS	<u>250MG</u>	<u>A070035 001</u>	Dec 20, 1984
<u>AB</u>	WATSON LABS INC	<u>500MG</u>	<u>A070044 001</u>	Feb 08, 1985
	INNOGENIX	125MG	A070772 003	Oct 29, 2024

METYRAPONE

CAPSULE; ORAL

METOPIRONE

+! HRA PHARMA 250MG N012911 002 Aug 09, 1996

METYROSINE

CAPSULE; ORAL

DEMSEERAB +! BAUSCH 250MG N017871 001METYROSINE

<u>AB</u>	AMNEAL	<u>250MG</u>	<u>A213734 001</u>	Jul 24, 2020
<u>AB</u>	LEADING	<u>250MG</u>	<u>A215541 001</u>	Sep 17, 2024

MEXILETINE HYDROCHLORIDE

CAPSULE; ORAL

MEXILETINE HYDROCHLORIDE

<u>AB</u>	ANI PHARMS	<u>150MG</u>	<u>A074450 001</u>	May 16, 1996
<u>AB</u>		<u>200MG</u>	<u>A074450 002</u>	May 16, 1996
<u>AB</u>		<u>250MG</u>	<u>A074450 003</u>	May 16, 1996
<u>AB</u>	ANNORA PHARMA	<u>150MG</u>	<u>A216463 001</u>	Nov 09, 2022
<u>AB</u>		<u>200MG</u>	<u>A216463 002</u>	Nov 09, 2022
<u>AB</u>		<u>250MG</u>	<u>A216463 003</u>	Nov 09, 2022
<u>AB</u>	CROSSMEDIKA SA	<u>150MG</u>	<u>A213500 001</u>	Jul 22, 2020
<u>AB</u>		<u>200MG</u>	<u>A213500 002</u>	Jul 22, 2020
<u>AB</u>		<u>250MG</u>	<u>A213500 003</u>	Jul 22, 2020
<u>AB</u>	LEADING	<u>150MG</u>	<u>A215876 001</u>	Feb 27, 2023
<u>AB</u>		<u>200MG</u>	<u>A215876 002</u>	Feb 27, 2023
<u>AB</u>		<u>250MG</u>	<u>A215876 003</u>	Feb 27, 2023
<u>AB</u>	NOVAST LABS	<u>150MG</u>	<u>A214352 001</u>	Jan 25, 2021
<u>AB</u>		<u>200MG</u>	<u>A214352 002</u>	Jan 25, 2021
<u>AB</u>		<u>250MG</u>	<u>A214352 003</u>	Jan 25, 2021
<u>AB</u>	RISING	<u>150MG</u>	<u>A215315 001</u>	Aug 26, 2022
<u>AB</u>		<u>200MG</u>	<u>A215315 002</u>	Aug 26, 2022
<u>AB</u>		<u>250MG</u>	<u>A215315 003</u>	Aug 26, 2022
<u>AB</u>	SENORES PHARMS	<u>150MG</u>	<u>A214089 001</u>	Oct 01, 2021
<u>AB</u>		<u>200MG</u>	<u>A214089 002</u>	Oct 01, 2021
<u>AB</u>		<u>250MG</u>	<u>A214089 003</u>	Oct 01, 2021
<u>AB</u>	TEVA	<u>150MG</u>	<u>A074377 001</u>	May 16, 1995
<u>AB</u>		<u>200MG</u>	<u>A074377 002</u>	May 16, 1995
<u>AB</u>	!	<u>250MG</u>	<u>A074377 003</u>	May 16, 1995

MICAFUNGIN SODIUM

INJECTABLE; INTRAVENOUS

MICAFUNGIN SODIUM

<u>AP</u>	BIOCON PHARMA	<u>EQ 50MG BASE/VIAL</u>	<u>A216438 001</u>	May 29, 2024
<u>AP</u>		<u>EQ 100MG BASE/VIAL</u>	<u>A216438 002</u>	May 29, 2024
<u>AP</u>	FRESENIUS KABI USA	<u>EQ 50MG BASE/VIAL</u>	<u>A207344 001</u>	May 17, 2019
<u>AP</u>		<u>EQ 100MG BASE/VIAL</u>	<u>A207344 002</u>	May 17, 2019
<u>AP</u>	HIKMA	<u>EQ 50MG BASE/VIAL</u>	<u>A213261 001</u>	Jul 09, 2021
<u>AP</u>		<u>EQ 100MG BASE/VIAL</u>	<u>A213261 002</u>	Jul 09, 2021
<u>AP</u>	JIANGSU HANSONH PHARM	<u>EQ 50MG BASE/VIAL</u>	<u>A213363 001</u>	Jul 09, 2021
<u>AP</u>		<u>EQ 100MG BASE/VIAL</u>	<u>A213363 002</u>	Jul 09, 2021
<u>AP</u>	MEITHEAL	<u>EQ 50MG BASE/VIAL</u>	<u>A215381 001</u>	Sep 28, 2022
<u>AP</u>		<u>EQ 100MG BASE/VIAL</u>	<u>A215381 002</u>	Sep 28, 2022
<u>AP</u>	XELLIA PHARMS APS	<u>EQ 50MG BASE/VIAL</u>	<u>A211713 001</u>	Jun 02, 2021
<u>AP</u>		<u>EQ 100MG BASE/VIAL</u>	<u>A211713 002</u>	Jun 02, 2021
<u>AP</u>	ZYDUS PHARMS	<u>EQ 50MG BASE/VIAL</u>	<u>A215241 001</u>	Oct 24, 2022

PRESCRIPTION DRUG PRODUCT LIST

MICAFUNGIN SODIUM

INJECTABLE; INTRAVENOUS

MICAFUNGIN SODIUM

AP		<u>EQ 100MG BASE/VIAL</u>	<u>A215241 002</u>	Oct 24, 2022
-----------	--	---------------------------	---------------------------	--------------

MYCAMINE

AP	+ !	ASTELLAS	<u>EQ 50MG BASE/VIAL</u>	<u>N021506 002</u>	Mar 16, 2005
-----------	------------	----------	--------------------------	---------------------------	--------------

AP	+ !		<u>EQ 100MG BASE/VIAL</u>	<u>N021506 003</u>	Jun 27, 2006
-----------	------------	--	---------------------------	---------------------------	--------------

POWDER; INTRAVENOUS

MICAFUNGIN

+ !	ENDO OPERATIONS	EQ 50MG BASE/VIAL	N212156 001	Jun 16, 2021
------------	-----------------	-------------------	-------------	--------------

+ !		EQ 100MG BASE/VIAL	N212156 002	Jun 16, 2021
------------	--	--------------------	-------------	--------------

SOLUTION; INTRAVENOUS

MICAFUNGIN IN SODIUM CHLORIDE 0.9%

+ !	BAXTER HLTHCARE CORP	EQ 50MG BASE/50ML (EQ 1MG BASE/ML)	N216142 001	Sep 29, 2023
------------	----------------------	------------------------------------	-------------	--------------

+ !		EQ 100MG BASE/100ML (EQ 1MG BASE/ML)	N216142 002	Sep 29, 2023
------------	--	--------------------------------------	-------------	--------------

+ !		EQ 150MG BASE/150ML (EQ 1MG BASE/ML)	N216142 003	Sep 29, 2023
------------	--	--------------------------------------	-------------	--------------

MICONAZOLE

TABLET; BUCCAL

ORAVIG

+ !	GALT PHARMS	50MG	N022404 001	Apr 16, 2010
------------	-------------	------	-------------	--------------

MICONAZOLE NITRATE

SUPPOSITORY; VAGINAL

MICONAZOLE NITRATE

AB	ACTAVIS PHARMA	<u>200MG</u>	<u>A073508 001</u>	Nov 19, 1993
-----------	----------------	---------------------	---------------------------	--------------

MONISTAT 3

AB	+ !	MEDTECH PRODUCTS	<u>200MG</u>	<u>N018888 001</u>	Aug 15, 1984
-----------	------------	------------------	---------------------	---------------------------	--------------

MICONAZOLE NITRATE; WHITE PETROLATUM; ZINC OXIDE

OINTMENT; TOPICAL

VUSION

+ !	MYLAN	0.25%; 81.35%; 15%	N021026 001	Feb 16, 2006
------------	-------	--------------------	-------------	--------------

MIDAZOLAM

SOLUTION; INTRAVENOUS

MIDAZOLAM IN 0.9% SODIUM CHLORIDE

AP	GLAND PHARMA LTD	<u>50MG/50ML (1MG/ML)</u>	<u>A218993 001</u>	Aug 12, 2024
-----------	------------------	---------------------------	---------------------------	--------------

AP		<u>100MG/100ML (1MG/ML)</u>	<u>A218993 002</u>	Aug 12, 2024
-----------	--	-----------------------------	---------------------------	--------------

AP	HIKMA	<u>50MG/50ML (1MG/ML)</u>	<u>A216159 001</u>	Apr 17, 2023
-----------	-------	---------------------------	---------------------------	--------------

AP		<u>100MG/100ML (1MG/ML)</u>	<u>A216159 002</u>	Apr 17, 2023
-----------	--	-----------------------------	---------------------------	--------------

AP	+ !	INFORLIFE	<u>50MG/50ML (1MG/ML)</u>	<u>N211844 001</u>	Mar 22, 2021
-----------	------------	-----------	---------------------------	---------------------------	--------------

AP	+ !		<u>100MG/100ML (1MG/ML)</u>	<u>N211844 002</u>	Mar 22, 2021
-----------	------------	--	-----------------------------	---------------------------	--------------

SPRAY; NASAL

NAYZILAM

+ !	UCB INC	5MG/SPRAY	N211321 001	May 17, 2019
------------	---------	-----------	-------------	--------------

MIDAZOLAM HYDROCHLORIDE

INJECTABLE; INJECTION

MIDAZOLAM HYDROCHLORIDE

AP	EPIC PHARMA LLC	<u>EQ 1MG BASE/ML</u>	<u>A075494 001</u>	Jun 30, 2000
-----------	-----------------	-----------------------	---------------------------	--------------

AP		<u>EQ 5MG BASE/ML</u>	<u>A075494 002</u>	Jun 30, 2000
-----------	--	-----------------------	---------------------------	--------------

AP	FRESENIUS KABI USA	<u>EQ 1MG BASE/ML</u>	<u>A075154 002</u>	Jun 20, 2000
-----------	--------------------	-----------------------	---------------------------	--------------

AP		<u>EQ 5MG BASE/ML</u>	<u>A075154 001</u>	Jun 20, 2000
-----------	--	-----------------------	---------------------------	--------------

AP	GLAND PHARMA LTD	<u>EQ 1MG BASE/ML</u>	<u>A090696 001</u>	Feb 29, 2012
-----------	------------------	-----------------------	---------------------------	--------------

AP		<u>EQ 5MG BASE/ML</u>	<u>A090850 001</u>	Jan 25, 2012
-----------	--	-----------------------	---------------------------	--------------

AP	HIKMA	<u>EQ 1MG BASE/ML</u>	<u>A075243 001</u>	Jun 20, 2000
-----------	-------	-----------------------	---------------------------	--------------

AP		<u>EQ 1MG BASE/ML</u>	<u>A075247 002</u>	Jun 23, 2000
-----------	--	-----------------------	---------------------------	--------------

AP		<u>EQ 1MG BASE/ML</u>	<u>A075324 001</u>	Jun 20, 2000
-----------	--	-----------------------	---------------------------	--------------

AP		<u>EQ 1MG BASE/ML</u>	<u>A075421 002</u>	Jun 20, 2000
-----------	--	-----------------------	---------------------------	--------------

AP		<u>EQ 1MG BASE/ML</u>	<u>A212847 001</u>	Dec 11, 2020
-----------	--	-----------------------	---------------------------	--------------

AP		<u>EQ 5MG BASE/ML</u>	<u>A075243 002</u>	Jun 20, 2000
-----------	--	-----------------------	---------------------------	--------------

AP		<u>EQ 5MG BASE/ML</u>	<u>A075247 001</u>	Jun 23, 2000
-----------	--	-----------------------	---------------------------	--------------

AP		<u>EQ 5MG BASE/ML</u>	<u>A075324 002</u>	Jun 20, 2000
-----------	--	-----------------------	---------------------------	--------------

AP		<u>EQ 5MG BASE/ML</u>	<u>A075421 001</u>	Jun 20, 2000
-----------	--	-----------------------	---------------------------	--------------

AP		<u>EQ 5MG BASE/ML</u>	<u>A212847 002</u>	Dec 11, 2020
-----------	--	-----------------------	---------------------------	--------------

AP	!	HOSPIRA	<u>EQ 1MG BASE/ML</u>	<u>A075293 001</u>	Jun 20, 2000
-----------	----------	---------	-----------------------	---------------------------	--------------

AP	!		<u>EQ 5MG BASE/ML</u>	<u>A075293 002</u>	Jun 20, 2000
-----------	----------	--	-----------------------	---------------------------	--------------

AP		MICRO LABS	<u>EQ 1MG BASE/ML</u>	<u>A217504 001</u>	Aug 21, 2023
-----------	--	------------	-----------------------	---------------------------	--------------

AP			<u>EQ 5MG BASE/ML</u>	<u>A217504 002</u>	Aug 21, 2023
-----------	--	--	-----------------------	---------------------------	--------------

MIDAZOLAM HYDROCHLORIDE PRESERVATIVE FREE

AP	FRESENIUS KABI USA	<u>EQ 1MG BASE/ML</u>	<u>A203460 001</u>	Aug 22, 2014
-----------	--------------------	-----------------------	---------------------------	--------------

AP		<u>EQ 5MG BASE/ML</u>	<u>A203460 002</u>	Aug 22, 2014
-----------	--	-----------------------	---------------------------	--------------

AP	!	HOSPIRA	<u>EQ 1MG BASE/ML</u>	<u>A075857 001</u>	Jul 22, 2002
-----------	----------	---------	-----------------------	---------------------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

MIDAZOLAM HYDROCHLORIDE

INJECTABLE; INJECTION

MIDAZOLAM HYDROCHLORIDE PRESERVATIVE FREE

<u>AP</u>	!		<u>EQ 5MG BASE/ML</u>	<u>A075857 002</u>	Jul 22, 2002
<u>AP</u>		STERISCIENCE SPECLTS	<u>EQ 1MG BASE/ML</u>	<u>A090315 001</u>	Nov 29, 2010
<u>AP</u>			<u>EQ 5MG BASE/ML</u>	<u>A090315 002</u>	Nov 29, 2010

MIDAZOLAM HYDROCHLORIDE

<u>AP</u>		STERISCIENCE SPECLTS	<u>EQ 1MG BASE/ML</u>	<u>A090316 001</u>	May 04, 2011
<u>AP</u>			<u>EQ 5MG BASE/ML</u>	<u>A090316 002</u>	May 04, 2011

SOLUTION; INTRAMUSCULAR

MIDAZOLAM HYDROCHLORIDE (AUTOINJECTOR)

+	!	RAFA LABS LTD	EQ 10MG BASE/0.7ML (EQ 10MG BASE/0.7ML)	N216359 001	Aug 08, 2022
---	---	---------------	---	-------------	--------------

SEIZALAM

+	!	MMT	EQ 50MG BASE/10ML (EQ 5MG BASE/ML)	N209566 001	Sep 14, 2018
---	---	-----	------------------------------------	-------------	--------------

SYRUP; ORAL

MIDAZOLAM HYDROCHLORIDE

<u>AA</u>	!	HIKMA	<u>EQ 2MG BASE/ML</u>	<u>A075873 001</u>	Apr 30, 2002
<u>AA</u>		PADAGIS US	<u>EQ 2MG BASE/ML</u>	<u>A076379 001</u>	May 02, 2005

MIDODRINE HYDROCHLORIDE

TABLET; ORAL

MIDODRINE HYDROCHLORIDE

<u>AB</u>		ALEMBIC	<u>2.5MG</u>	<u>A214734 001</u>	Jan 21, 2021
<u>AB</u>			<u>5MG</u>	<u>A214734 002</u>	Jan 21, 2021
<u>AB</u>			<u>10MG</u>	<u>A214734 003</u>	Jan 21, 2021
<u>AB</u>		APOTEX	<u>2.5MG</u>	<u>A077746 001</u>	Sep 12, 2006
<u>AB</u>			<u>5MG</u>	<u>A077746 002</u>	Sep 12, 2006
<u>AB</u>			<u>10MG</u>	<u>A077746 003</u>	Sep 12, 2006
<u>AB</u>		AUROBINDO PHARMA LTD	<u>2.5MG</u>	<u>A212774 001</u>	Aug 10, 2020
<u>AB</u>			<u>5MG</u>	<u>A212774 002</u>	Aug 10, 2020
<u>AB</u>			<u>10MG</u>	<u>A212774 003</u>	Aug 10, 2020
<u>AB</u>		ENDO OPERATIONS	<u>2.5MG</u>	<u>A207169 001</u>	Oct 29, 2018
<u>AB</u>			<u>5MG</u>	<u>A207169 002</u>	Oct 29, 2018
<u>AB</u>			<u>10MG</u>	<u>A207169 003</u>	Oct 29, 2018
<u>AB</u>		IMPAX PHARMS	<u>2.5MG</u>	<u>A076449 001</u>	May 27, 2004
<u>AB</u>			<u>5MG</u>	<u>A076449 002</u>	May 27, 2004
<u>AB</u>			<u>10MG</u>	<u>A076449 003</u>	Dec 16, 2005
<u>AB</u>		MANKIND PHARMA	<u>2.5MG</u>	<u>A217271 001</u>	Nov 01, 2023
<u>AB</u>			<u>5MG</u>	<u>A217271 002</u>	Nov 01, 2023
<u>AB</u>			<u>10MG</u>	<u>A217271 003</u>	Nov 01, 2023
<u>AB</u>		MYLAN PHARMS INC	<u>2.5MG</u>	<u>A076577 001</u>	Sep 10, 2003
<u>AB</u>			<u>5MG</u>	<u>A076577 002</u>	Sep 10, 2003
<u>AB</u>			<u>10MG</u>	<u>A076577 003</u>	Sep 10, 2003
<u>AB</u>		NOVUGEN	<u>2.5MG</u>	<u>A211973 001</u>	Oct 17, 2023
<u>AB</u>			<u>5MG</u>	<u>A211973 002</u>	Oct 17, 2023
<u>AB</u>			<u>10MG</u>	<u>A211973 003</u>	Oct 17, 2023
<u>AB</u>		RUBICON	<u>2.5MG</u>	<u>A212543 001</u>	Aug 19, 2019
<u>AB</u>			<u>5MG</u>	<u>A212543 002</u>	Aug 19, 2019
<u>AB</u>			<u>10MG</u>	<u>A212543 003</u>	Aug 19, 2019
<u>AB</u>		THINQ PHARM-CRO PVT	<u>2.5MG</u>	<u>A207613 001</u>	Nov 02, 2018
<u>AB</u>			<u>5MG</u>	<u>A207613 002</u>	Nov 02, 2018
<u>AB</u>			<u>10MG</u>	<u>A207613 003</u>	Nov 02, 2018
<u>AB</u>		XIROMED	<u>2.5MG</u>	<u>A207849 001</u>	Oct 01, 2020
<u>AB</u>			<u>5MG</u>	<u>A207849 002</u>	Oct 01, 2020
<u>AB</u>			<u>10MG</u>	<u>A207849 003</u>	Oct 01, 2020
<u>AB</u>		ZYDUS LIFESCIENCES	<u>2.5MG</u>	<u>A213055 001</u>	Sep 01, 2020
<u>AB</u>			<u>5MG</u>	<u>A213055 002</u>	Sep 01, 2020
<u>AB</u>			<u>10MG</u>	<u>A213055 003</u>	Sep 01, 2020

ORVATEN

<u>AB</u>		UPSHER SMITH LABS	<u>2.5MG</u>	<u>A076725 001</u>	Nov 03, 2004
<u>AB</u>	!		<u>5MG</u>	<u>A076725 002</u>	Nov 03, 2004
<u>AB</u>			<u>10MG</u>	<u>A076725 003</u>	Nov 03, 2004

MIDOSTAURIN

CAPSULE; ORAL

MIDOSTAURIN

<u>AB</u>		DR REDDYS	<u>25MG</u>	<u>A215921 001</u>	Jun 28, 2024
<u>AB</u>		LUPIN	<u>25MG</u>	<u>A216015 001</u>	May 10, 2024

RYDAPT

<u>AB</u>	+	NOVARTIS	<u>25MG</u>	<u>N207997 001</u>	Apr 28, 2017
-----------	---	----------	-------------	--------------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

MIFEPRISTONE

TABLET; ORAL

KORLYM

AB	+ !	CORCEPT THERAP	300MG	N202107	001	Feb 17, 2012
-----------	------------	----------------	--------------	----------------	------------	--------------

MIFEPREX

AB	+ !	DANCO LABS LLC	200MG	N020687	001	Sep 28, 2000
-----------	------------	----------------	--------------	----------------	------------	--------------

MIFEPRISTONE

AB		GENBIOPRO	200MG	A091178	001	Apr 11, 2019
-----------	--	-----------	--------------	----------------	------------	--------------

AB		TEVA PHARMS USA INC	300MG	A211436	001	Aug 03, 2020
-----------	--	---------------------	--------------	----------------	------------	--------------

MIGALASTAT HYDROCHLORIDE

CAPSULE; ORAL

GALAFOLD

+ !	AMICUS THERAP US	EQ 123MG BASE	N208623	001	Aug 10, 2018
------------	------------------	---------------	---------	-----	--------------

MIGLITOL

TABLET; ORAL

GLYSET

AB	+ !	PFIZER	25MG	N020682	001	Dec 18, 1996
-----------	------------	--------	-------------	----------------	------------	--------------

AB	+		50MG	N020682	002	Dec 18, 1996
-----------	----------	--	-------------	----------------	------------	--------------

AB	+		100MG	N020682	003	Dec 18, 1996
-----------	----------	--	--------------	----------------	------------	--------------

MIGLITOL

AB		WESTMINSTER PHARMS	25MG	A203965	001	Feb 24, 2015
-----------	--	--------------------	-------------	----------------	------------	--------------

AB			50MG	A203965	002	Feb 24, 2015
-----------	--	--	-------------	----------------	------------	--------------

AB			100MG	A203965	003	Feb 24, 2015
-----------	--	--	--------------	----------------	------------	--------------

MIGLUSTAT

CAPSULE; ORAL

MIGLUSTAT

AB		ANI PHARMS	100MG	A208342	001	Apr 17, 2018
-----------	--	------------	--------------	----------------	------------	--------------

YARGESA

AB		EDENBRIDGE PHARMS	100MG	A209821	001	Aug 06, 2020
-----------	--	-------------------	--------------	----------------	------------	--------------

ZAVESCA

AB	+ !	ACTELION	100MG	N021348	001	Jul 31, 2003
-----------	------------	----------	--------------	----------------	------------	--------------

OPFOLDA

+ !	AMICUS THERAP US	65MG	N215211	001	Sep 28, 2023
------------	------------------	------	---------	-----	--------------

MILNACIPRAN HYDROCHLORIDE

TABLET; ORAL

MILNACIPRAN HYDROCHLORIDE

AB		HETERO LABS LTD V	12.5MG	A205147	001	Oct 03, 2024
-----------	--	-------------------	---------------	----------------	------------	--------------

AB			25MG	A205147	002	Oct 03, 2024
-----------	--	--	-------------	----------------	------------	--------------

AB			50MG	A205147	003	Oct 03, 2024
-----------	--	--	-------------	----------------	------------	--------------

AB			100MG	A205147	004	Oct 03, 2024
-----------	--	--	--------------	----------------	------------	--------------

SAVELLA

AB	+	ABBVIE	12.5MG	N022256	001	Jan 14, 2009
-----------	----------	--------	---------------	----------------	------------	--------------

AB	+		25MG	N022256	002	Jan 14, 2009
-----------	----------	--	-------------	----------------	------------	--------------

AB	+ !		50MG	N022256	003	Jan 14, 2009
-----------	------------	--	-------------	----------------	------------	--------------

AB	+		100MG	N022256	004	Jan 14, 2009
-----------	----------	--	--------------	----------------	------------	--------------

MILRINONE LACTATE

INJECTABLE; INJECTION

MILRINONE LACTATE

AP		CAPLIN	EQ 1MG BASE/ML	A214380	001	Apr 16, 2021
-----------	--	--------	-----------------------	----------------	------------	--------------

AP		FRESENIUS KABI USA	EQ 1MG BASE/ML	A075936	001	May 28, 2002
-----------	--	--------------------	-----------------------	----------------	------------	--------------

AP		HIKMA	EQ 1MG BASE/ML	A075530	001	May 28, 2002
-----------	--	-------	-----------------------	----------------	------------	--------------

AP			EQ 1MG BASE/ML	A075660	001	May 28, 2002
-----------	--	--	-----------------------	----------------	------------	--------------

AP	!	HIKMA FARMACEUTICA	EQ 1MG BASE/ML	A077966	001	Dec 03, 2010
-----------	----------	--------------------	-----------------------	----------------	------------	--------------

AP		HOSPIRA	EQ 1MG BASE/ML	A203280	001	Sep 03, 2014
-----------	--	---------	-----------------------	----------------	------------	--------------

AP		MEITHEAL	EQ 1MG BASE/ML	A211671	001	Mar 24, 2020
-----------	--	----------	-----------------------	----------------	------------	--------------

AP		SHANDONG	EQ 1MG BASE/ML	A216373	001	Jan 23, 2023
-----------	--	----------	-----------------------	----------------	------------	--------------

MILRINONE LACTATE IN DEXTROSE 5%

AP		EUGIA PHARMA	EQ 20MG BASE/100ML (EQ 0.2MG BASE/ML)	A209666	001	Sep 03, 2020
-----------	--	--------------	--	----------------	------------	--------------

AP			EQ 40MG BASE/200ML (EQ 0.2MG BASE/ML)	A209666	002	Sep 03, 2020
-----------	--	--	--	----------------	------------	--------------

MILRINONE LACTATE IN DEXTROSE 5% IN PLASTIC CONTAINER

AP	!	BAXTER HLTHCARE	EQ 20MG BASE/100ML (EQ 0.2MG BASE/ML)	A075834	001	May 28, 2002
-----------	----------	-----------------	--	----------------	------------	--------------

AP	!		EQ 40MG BASE/200ML (EQ 0.2MG BASE/ML)	A075834	002	May 28, 2002
-----------	----------	--	--	----------------	------------	--------------

AP		GLAND PHARMA LTD	EQ 20MG BASE/100ML (EQ 0.2MG BASE/ML)	A213585	001	Jul 16, 2020
-----------	--	------------------	--	----------------	------------	--------------

AP			EQ 40MG BASE/200ML (EQ 0.2MG BASE/ML)	A213585	002	Jul 16, 2020
-----------	--	--	--	----------------	------------	--------------

AP		HIKMA	EQ 20MG BASE/100ML (EQ 0.2MG BASE/ML)	A078113	001	May 21, 2008
-----------	--	-------	--	----------------	------------	--------------

AP			EQ 40MG BASE/200ML (EQ 0.2MG BASE/ML)	A078113	002	May 21, 2008
-----------	--	--	--	----------------	------------	--------------

MILRINONE LACTATE IN PLASTIC CONTAINER

AP		HIKMA FARMACEUTICA	EQ 20MG BASE/100ML (EQ 0.2MG BASE/ML)	A090038	001	Jan 21, 2010
-----------	--	--------------------	--	----------------	------------	--------------

AP			EQ 40MG BASE/200ML (EQ 0.2MG BASE/ML)	A090038	002	Jan 21, 2010
-----------	--	--	--	----------------	------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

MILTEFOSINE

CAPSULE;ORAL

IMPAVIDO

+! KNIGHT THERAPS 50MG N204684 001 Mar 19, 2014

MINOCYCLINE HYDROCHLORIDE

AEROSOL, FOAM;TOPICAL

AMZEEQ

+! JOURNEY EQ 4% BASE N212379 001 Oct 18, 2019

ZILXI

+! JOURNEY EQ 1.5% BASE N213690 001 May 28, 2020

CAPSULE;ORAL

MINOCINAB + BAUSCH EQ 50MG BASE N050649 001 May 31, 1990AB + EQ 100MG BASE N050649 002 May 31, 1990MINOCYCLINE HYDROCHLORIDEAB AUROBINDO PHARMA EQ 50MG BASE A065470 001 Mar 11, 2008AB EQ 75MG BASE A065470 002 Mar 11, 2008AB EQ 100MG BASE A065470 003 Mar 11, 2008AB IMPAX LABS EQ 50MG BASE A065005 001 Mar 23, 1999AB EQ 75MG BASE A065005 003 Apr 18, 2001AB EQ 100MG BASE A065005 002 Mar 23, 1999AB SUN PHARM INDS INC EQ 50MG BASE A090867 001 May 13, 2013AB EQ 75MG BASE A090867 002 May 13, 2013AB EQ 100MG BASE A090867 003 May 13, 2013AB TORRENT EQ 50MG BASE A065062 001 Nov 30, 2000AB EQ 75MG BASE A065062 002 Nov 30, 2000AB EQ 100MG BASE A065062 003 Nov 30, 2000AB WATSON LABS EQ 75MG BASE A063065 002 Jun 10, 1999AB EQ 100MG BASE A063065 001 Dec 30, 1991AB WATSON LABS TEVA EQ 50MG BASE A063181 001 Dec 30, 1991AB ZYDUS EQ 50MG BASE A063011 001 Mar 02, 1992AB EQ 75MG BASE A063009 002 Aug 12, 2003AB ! EQ 100MG BASE A063009 001 Mar 02, 1992

CAPSULE, EXTENDED RELEASE;ORAL

EMROSI

+! JOURNEY 40MG N219015 001 Nov 01, 2024

INJECTABLE;INJECTION

MINOCINAP +! REMPEX EQ 100MG BASE/VIAL N050444 001MINOCYCLINE HYDROCHLORIDEAP NEXUS PHARMS EQ 100MG BASE/VIAL A214934 001 Oct 24, 2024

POWDER, EXTENDED RELEASE;DENTAL

ARESTIN

+! ORAPHARMA EQ 1MG BASE N050781 001 Feb 16, 2001

TABLET;ORAL

MINOCYCLINE HYDROCHLORIDEAB AUROBINDO PHARMA EQ 50MG BASE A213662 001 May 01, 2020AB LTD EQ 75MG BASE A213662 002 May 01, 2020AB EQ 100MG BASE A213662 003 May 01, 2020AB BEXIMCO PHARMS USA EQ 50MG BASE A215466 001 May 27, 2022AB EQ 75MG BASE A215466 002 May 27, 2022AB EQ 100MG BASE A215466 003 May 27, 2022AB CHARTWELL RX EQ 50MG BASE A065436 001 Dec 26, 2007AB EQ 75MG BASE A065436 002 Dec 26, 2007AB EQ 100MG BASE A065436 003 Dec 26, 2007AB STRIDES PHARMA EQ 50MG BASE A065131 001 Apr 16, 2003AB EQ 75MG BASE A065131 002 Apr 16, 2003AB ! EQ 100MG BASE A065131 003 Apr 16, 2003AB TORRENT EQ 50MG BASE A065156 001 Jan 06, 2004AB EQ 75MG BASE A065156 002 Jan 06, 2004AB EQ 100MG BASE A065156 003 Jan 06, 2004

TABLET, EXTENDED RELEASE;ORAL

MINOCYCLINE HYDROCHLORIDEAB ALKEM LABS LTD EQ 45MG BASE A204453 001 Sep 28, 2016AB EQ 55MG BASE A204453 008 Dec 19, 2019AB EQ 65MG BASE A204453 006 Mar 16, 2018AB EQ 80MG BASE A204453 002 Sep 28, 2016AB EQ 90MG BASE A204453 003 Sep 28, 2016AB EQ 105MG BASE A204453 004 Sep 28, 2016AB ! EQ 115MG BASE A204453 007 Mar 16, 2018AB ! EQ 135MG BASE A204453 005 Sep 28, 2016

PRESCRIPTION DRUG PRODUCT LIST

MINOCYCLINE HYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL

MINOCYCLINE HYDROCHLORIDE

<u>AB</u>	AUROBINDO PHARMA LTD	<u>EQ 45MG BASE</u>	<u>A202261 001</u>	Nov 19, 2012
<u>AB</u>		<u>EQ 55MG BASE</u>	<u>A202261 008</u>	Aug 21, 2019
<u>AB</u>		<u>EQ 65MG BASE</u>	<u>A202261 002</u>	Sep 28, 2018
<u>AB</u>		<u>EQ 80MG BASE</u>	<u>A202261 006</u>	Jun 13, 2016
<u>AB</u>		<u>EQ 90MG BASE</u>	<u>A202261 003</u>	Nov 19, 2012
<u>AB</u>		<u>EQ 105MG BASE</u>	<u>A202261 007</u>	Jun 13, 2016
<u>AB</u>		<u>EQ 115MG BASE</u>	<u>A202261 004</u>	Sep 28, 2018
<u>AB</u>		<u>EQ 135MG BASE</u>	<u>A202261 005</u>	Nov 19, 2012
<u>AB</u>	CHARTWELL RX	<u>EQ 45MG BASE</u>	<u>A091424 001</u>	Nov 30, 2011
<u>AB</u>		<u>EQ 55MG BASE</u>	<u>A091424 002</u>	Nov 30, 2011
<u>AB</u>		<u>EQ 90MG BASE</u>	<u>A091424 003</u>	Nov 30, 2011
<u>AB</u>		<u>EQ 135MG BASE</u>	<u>A091424 004</u>	Nov 30, 2011
<u>AB</u>	SANDOZ	<u>EQ 45MG BASE</u>	<u>A090422 001</u>	Aug 13, 2009
<u>AB</u>		<u>EQ 90MG BASE</u>	<u>A090422 002</u>	Aug 13, 2009
<u>AB</u>		<u>EQ 135MG BASE</u>	<u>A090422 003</u>	Aug 13, 2009
<u>AB</u>	SUN PHARM	<u>EQ 45MG BASE</u>	<u>A204394 001</u>	Dec 30, 2015
<u>AB</u>		<u>EQ 55MG BASE</u>	<u>A204394 002</u>	Oct 07, 2022
<u>AB</u>		<u>EQ 65MG BASE</u>	<u>A204394 003</u>	Oct 07, 2022
<u>AB</u>		<u>EQ 80MG BASE</u>	<u>A204394 004</u>	Dec 30, 2015
<u>AB</u>		<u>EQ 105MG BASE</u>	<u>A204394 005</u>	Dec 30, 2015
<u>AB</u>		<u>EQ 115MG BASE</u>	<u>A204394 006</u>	Oct 07, 2022
<u>AB</u>		<u>EQ 135MG BASE</u>	<u>A204394 007</u>	Dec 30, 2015

MINOXIDIL

TABLET;ORAL

MINOXIDIL

<u>AB</u>	ENDO OPERATIONS	<u>2.5MG</u>	<u>A071826 001</u>	Nov 14, 1988
<u>AB</u>		<u>10MG</u>	<u>A071839 001</u>	Nov 14, 1988
<u>AB</u>	SUN PHARM INDUSTRIES	<u>2.5MG</u>	<u>A072709 002</u>	Dec 14, 1995
<u>AB</u>		<u>10MG</u>	<u>A072709 001</u>	Dec 14, 1995
<u>AB</u>	WATSON LABS	<u>2.5MG</u>	<u>A071344 001</u>	Mar 03, 1987
<u>AB</u>	!	<u>10MG</u>	<u>A071345 001</u>	Mar 03, 1987

MIRABEGRON

FOR SUSPENSION, EXTENDED RELEASE;ORAL

MYRBETRIQ GRANULES

+! APGDI 8MG/ML

N213801 001 Mar 25, 2021

TABLET, EXTENDED RELEASE;ORAL

MIRABEGRON

<u>AB</u>	ALKEM LABS LTD	<u>25MG</u>	<u>A215948 001</u>	Feb 12, 2024
<u>AB</u>		<u>50MG</u>	<u>A215948 002</u>	Feb 12, 2024
<u>AB</u>	APOTEX	<u>25MG</u>	<u>A209434 001</u>	Jan 02, 2025
<u>AB</u>		<u>50MG</u>	<u>A209434 002</u>	Jan 02, 2025
<u>AB</u>	LUPIN LTD	<u>25MG</u>	<u>A209485 001</u>	Sep 28, 2022
<u>AB</u>		<u>50MG</u>	<u>A209485 002</u>	Sep 28, 2022
<u>AB</u>	ZYDUS PHARMS	<u>25MG</u>	<u>A209488 001</u>	Sep 29, 2022
<u>AB</u>		<u>50MG</u>	<u>A209488 002</u>	Sep 29, 2022
	<u>MYRBETRIQ</u>			
<u>AB</u>	+! APGDI	<u>25MG</u>	<u>N202611 001</u>	Jun 28, 2012
<u>AB</u>	+!	<u>50MG</u>	<u>N202611 002</u>	Jun 28, 2012

MIRTAZAPINE

TABLET;ORAL

MIRTAZAPINE

<u>AB</u>	APOTEX INC	<u>15MG</u>	<u>A077666 001</u>	Aug 22, 2007
<u>AB</u>		<u>30MG</u>	<u>A077666 002</u>	Aug 22, 2007
<u>AB</u>		<u>45MG</u>	<u>A077666 003</u>	Aug 22, 2007
<u>AB</u>	AUROBINDO	<u>7.5MG</u>	<u>A076921 001</u>	Oct 22, 2004
<u>AB</u>		<u>15MG</u>	<u>A076921 002</u>	Oct 22, 2004
<u>AB</u>		<u>30MG</u>	<u>A076921 003</u>	Oct 22, 2004
<u>AB</u>		<u>45MG</u>	<u>A076921 004</u>	Oct 22, 2004
<u>AB</u>	CHARTWELL RX	<u>15MG</u>	<u>A076219 001</u>	Jun 19, 2003
<u>AB</u>		<u>30MG</u>	<u>A076219 002</u>	Jun 19, 2003
<u>AB</u>		<u>45MG</u>	<u>A076219 003</u>	Jun 19, 2003
<u>AB</u>	MYLAN	<u>15MG</u>	<u>A076122 001</u>	Jun 19, 2003
<u>AB</u>		<u>30MG</u>	<u>A076122 002</u>	Jun 19, 2003
<u>AB</u>		<u>45MG</u>	<u>A076122 003</u>	Jun 19, 2003
<u>AB</u>	PRASCO	<u>7.5MG</u>	<u>A216751 001</u>	Jan 18, 2023
<u>AB</u>		<u>15MG</u>	<u>A216751 002</u>	Jan 18, 2023

PRESCRIPTION DRUG PRODUCT LIST

MIRTAZAPINE

TABLET; ORAL

MIRTAZAPINE

<u>AB</u>		<u>30MG</u>	<u>A216751 003</u>	Jan 18, 2023	
<u>AB</u>		<u>45MG</u>	<u>A216751 004</u>	Jan 18, 2023	
<u>AB</u>	!	SUN PHARM INDS INC	<u>7.5MG</u>	<u>A076541 004</u>	Apr 22, 2004
<u>AB</u>			<u>15MG</u>	<u>A076541 001</u>	Apr 22, 2004
<u>AB</u>			<u>30MG</u>	<u>A076541 002</u>	Apr 22, 2004
<u>AB</u>			<u>45MG</u>	<u>A076541 003</u>	Apr 22, 2004
<u>AB</u>		TEVA	<u>15MG</u>	<u>A076119 001</u>	Jan 24, 2003
<u>AB</u>			<u>30MG</u>	<u>A076119 002</u>	Jan 24, 2003
<u>AB</u>			<u>45MG</u>	<u>A076119 003</u>	Jun 19, 2003

REMERON

<u>AB</u>	+	!	ORGANON	<u>15MG</u>	<u>N020415 001</u>	Jun 14, 1996
<u>AB</u>	+			<u>30MG</u>	<u>N020415 002</u>	Jun 14, 1996

TABLET, ORALLY DISINTEGRATING; ORAL

MIRTAZAPINE

<u>AB</u>		AUROBINDO PHARMA	<u>15MG</u>	<u>A077376 002</u>	Dec 08, 2005
<u>AB</u>			<u>30MG</u>	<u>A077376 003</u>	Dec 08, 2005
<u>AB</u>			<u>45MG</u>	<u>A077376 004</u>	Feb 28, 2006
<u>AB</u>		SQUARE PHARMS	<u>15MG</u>	<u>A205798 001</u>	Jun 01, 2017
<u>AB</u>			<u>30MG</u>	<u>A205798 002</u>	Jun 01, 2017
<u>AB</u>			<u>45MG</u>	<u>A205798 003</u>	Jun 01, 2017

REMERON SOLTAB

<u>AB</u>	+	!	ORGANON USA ORGANON	<u>15MG</u>	<u>N021208 001</u>	Jan 12, 2001
<u>AB</u>	+			<u>30MG</u>	<u>N021208 002</u>	Jan 12, 2001
<u>AB</u>	+			<u>45MG</u>	<u>N021208 003</u>	Jan 12, 2001

MISOPROSTOL

TABLET; ORAL

CYTOTEC

<u>AB</u>	+		PFIZER	<u>0.1MG</u>	<u>N019268 003</u>	Sep 21, 1990
<u>AB</u>	+	!		<u>0.2MG</u>	<u>N019268 001</u>	Dec 27, 1988

MISOPROSTOL

<u>AB</u>		ACQ PHARMA	<u>0.1MG</u>	<u>A210201 001</u>	Jul 02, 2019
<u>AB</u>			<u>0.2MG</u>	<u>A210201 002</u>	Jul 02, 2019
<u>AB</u>		ANI PHARMS	<u>0.1MG</u>	<u>A076095 001</u>	Jul 10, 2002
<u>AB</u>			<u>0.2MG</u>	<u>A076095 002</u>	Jul 10, 2002
<u>AB</u>		MICRO LABS	<u>0.1MG</u>	<u>A216872 001</u>	Oct 17, 2022
<u>AB</u>			<u>0.2MG</u>	<u>A216872 002</u>	Oct 17, 2022

MITAPIVAT SULFATE

TABLET; ORAL

PYRUKYND

	+		AGIOS PHARMS INC	EQ 5MG BASE	N216196 001	Feb 17, 2022
	+			EQ 20MG BASE	N216196 002	Feb 17, 2022
	+	!		EQ 50MG BASE	N216196 003	Feb 17, 2022

MITOMYCIN

FOR SOLUTION; TOPICAL

MITOSOL

	+	!	MOBIUS THERAP	0.2MG/VIAL	N022572 001	Feb 07, 2012
--	---	---	---------------	------------	-------------	--------------

INJECTABLE; INJECTION

MITOMYCIN

<u>AP</u>	!		ACCORD HLTHCARE	<u>5MG/VIAL</u>	<u>A064144 001</u>	Apr 30, 1998
<u>AP</u>	!			<u>20MG/VIAL</u>	<u>A064144 002</u>	Apr 30, 1998
<u>AP</u>	!			<u>40MG/VIAL</u>	<u>A064144 003</u>	Aug 11, 2009
<u>AP</u>			EUGIA PHARMA	<u>5MG/VIAL</u>	<u>A216732 001</u>	Oct 30, 2023
<u>AP</u>				<u>20MG/VIAL</u>	<u>A216732 002</u>	Oct 30, 2023
<u>AP</u>				<u>40MG/VIAL</u>	<u>A216732 003</u>	Oct 30, 2023
<u>AP</u>			FRESENIUS KABI USA	<u>5MG/VIAL</u>	<u>A211269 001</u>	Apr 05, 2024
<u>AP</u>				<u>20MG/VIAL</u>	<u>A211269 002</u>	Apr 05, 2024
<u>AP</u>				<u>40MG/VIAL</u>	<u>A211269 003</u>	Apr 05, 2024
<u>AP</u>			GLAND PHARMA LTD	<u>5MG/VIAL</u>	<u>A215687 001</u>	Oct 20, 2021
<u>AP</u>				<u>20MG/VIAL</u>	<u>A215687 002</u>	Oct 20, 2021
<u>AP</u>				<u>40MG/VIAL</u>	<u>A216648 001</u>	Nov 22, 2022
<u>AP</u>			HIKMA	<u>5MG/VIAL</u>	<u>A064180 001</u>	Dec 23, 1999
<u>AP</u>				<u>20MG/VIAL</u>	<u>A064117 002</u>	Apr 19, 1995
<u>AP</u>				<u>20MG/VIAL</u>	<u>A064180 002</u>	Dec 23, 1999
<u>AP</u>				<u>40MG/VIAL</u>	<u>A064117 003</u>	Jun 02, 1999
<u>AP</u>			MEITHEAL	<u>5MG/VIAL</u>	<u>A214505 001</u>	Sep 08, 2022
<u>AP</u>				<u>20MG/VIAL</u>	<u>A214505 002</u>	Sep 08, 2022
<u>AP</u>				<u>40MG/VIAL</u>	<u>A214504 001</u>	Sep 06, 2022
<u>AP</u>			RK PHARMA	<u>5MG/VIAL</u>	<u>A202670 001</u>	Oct 13, 2017

PRESCRIPTION DRUG PRODUCT LIST

MITOMYCIN

INJECTABLE; INJECTION

MITOMYCIN

<u>AP</u>		<u>20MG/VIAL</u>	<u>A202670 002</u>	Oct 13, 2017
<u>AP</u>		<u>40MG/VIAL</u>	<u>A203386 001</u>	Oct 13, 2017
	POWDER; PYELOCALYCEAL JELMYTO			
	+! UROGEN PHARMA	40MG/VIAL	N211728 001	Apr 15, 2020

MITOTANE

TABLET; ORAL

LYSODREN

+!	HRA PHARMA	500MG	N016885 001	
----	------------	-------	-------------	--

MITOXANTRONE HYDROCHLORIDE

INJECTABLE; INJECTION

MITOXANTRONE HYDROCHLORIDE

<u>AP</u>	FRESENIUS KABI USA	<u>EQ 20MG BASE/10ML (EQ 2MG BASE/ML)</u>	<u>A077496 001</u>	Apr 11, 2006
<u>AP</u>		<u>EQ 25MG BASE/12.5ML (EQ 2MG BASE/ML)</u>	<u>A077496 002</u>	Apr 11, 2006
<u>AP</u>		<u>EQ 30MG BASE/15ML (EQ 2MG BASE/ML)</u>	<u>A077496 003</u>	Apr 11, 2006
<u>AP</u>	HIKMA	<u>EQ 20MG BASE/10ML (EQ 2MG BASE/ML)</u>	<u>A076611 001</u>	Apr 11, 2006
<u>AP</u>		<u>EQ 25MG BASE/12.5ML (EQ 2MG BASE/ML)</u>	<u>A076611 002</u>	Apr 11, 2006
<u>AP</u>		<u>EQ 30MG BASE/15ML (EQ 2MG BASE/ML)</u>	<u>A076611 003</u>	Apr 11, 2006
<u>AP</u>	! HOSPIRA	<u>EQ 20MG BASE/10ML (EQ 2MG BASE/ML)</u>	<u>A076871 001</u>	Apr 11, 2006
<u>AP</u>	!	<u>EQ 25MG BASE/12.5ML (EQ 2MG BASE/ML)</u>	<u>A076871 002</u>	Apr 11, 2006
<u>AP</u>	!	<u>EQ 30MG BASE/15ML (EQ 2MG BASE/ML)</u>	<u>A076871 003</u>	Apr 11, 2006
<u>AP</u>	MEITHEAL	<u>EQ 20MG BASE/10ML (EQ 2MG BASE/ML)</u>	<u>A077356 001</u>	Apr 11, 2006
<u>AP</u>		<u>EQ 25MG BASE/12.5ML (EQ 2MG BASE/ML)</u>	<u>A077356 002</u>	Apr 11, 2006
<u>AP</u>		<u>EQ 30MG BASE/15ML (EQ 2MG BASE/ML)</u>	<u>A077356 003</u>	Apr 11, 2006

MODAFINIL

TABLET; ORAL

MODAFINIL

<u>AB</u>	ALEMBIC	<u>100MG</u>	<u>A202700 001</u>	Oct 18, 2012
<u>AB</u>		<u>200MG</u>	<u>A202700 002</u>	Oct 18, 2012
<u>AB</u>	APOTEX	<u>100MG</u>	<u>A077667 001</u>	Feb 03, 2014
<u>AB</u>		<u>200MG</u>	<u>A077667 002</u>	Feb 03, 2014
<u>AB</u>	APPCO	<u>100MG</u>	<u>A207196 001</u>	Aug 16, 2017
<u>AB</u>		<u>200MG</u>	<u>A207196 002</u>	Aug 16, 2017
<u>AB</u>	AUROBINDO PHARMA LTD	<u>100MG</u>	<u>A202566 001</u>	Sep 27, 2012
<u>AB</u>		<u>200MG</u>	<u>A202566 002</u>	Sep 27, 2012
<u>AB</u>	CADILA	<u>100MG</u>	<u>A209966 001</u>	Sep 14, 2017
<u>AB</u>		<u>200MG</u>	<u>A209966 002</u>	Sep 14, 2017
<u>AB</u>	ORBION PHARMS	<u>100MG</u>	<u>A078963 001</u>	Sep 26, 2012
<u>AB</u>		<u>200MG</u>	<u>A078963 002</u>	Sep 26, 2012
<u>AB</u>	WATSON LABS INC	<u>100MG</u>	<u>A076715 001</u>	Nov 01, 2012
<u>AB</u>		<u>200MG</u>	<u>A076715 002</u>	Nov 01, 2012
	<u>PROVIGIL</u>			
<u>AB</u>	+ CEPHALON	<u>100MG</u>	<u>N020717 001</u>	Dec 24, 1998
<u>AB</u>	+!	<u>200MG</u>	<u>N020717 002</u>	Dec 24, 1998

MOEXIPRIL HYDROCHLORIDE

TABLET; ORAL

MOEXIPRIL HYDROCHLORIDE

<u>AB</u>	APOTEX INC	<u>7.5MG</u>	<u>A078454 001</u>	Jun 02, 2008
<u>AB</u>		<u>15MG</u>	<u>A078454 002</u>	Jun 02, 2008
<u>AB</u>	CHARTWELL RX	<u>7.5MG</u>	<u>A077536 001</u>	Nov 30, 2006
<u>AB</u>		<u>15MG</u>	<u>A077536 002</u>	Nov 30, 2006
<u>AB</u>	GLENMARK PHARMS LTD	<u>7.5MG</u>	<u>A090416 001</u>	Mar 30, 2010
<u>AB</u>		<u>15MG</u>	<u>A090416 002</u>	Mar 30, 2010
<u>AB</u>	TEVA	<u>7.5MG</u>	<u>A076204 001</u>	May 08, 2003
<u>AB</u>	!	<u>15MG</u>	<u>A076204 002</u>	May 08, 2003

MOLINDONE HYDROCHLORIDE

TABLET; ORAL

MOLINDONE HYDROCHLORIDE

	EPIC PHARMA LLC	5MG	A090453 001	Mar 20, 2015
		10MG	A090453 002	Mar 20, 2015
!		25MG	A090453 003	Mar 20, 2015

PRESCRIPTION DRUG PRODUCT LIST

MOMELOTINIB DIHYDROCHLORIDE

TABLET; ORAL

OJJAARA

+	GLAXOSMITHKLINE	EQ 100MG BASE	N216873	001	Sep 15, 2023
+		EQ 150MG BASE	N216873	002	Sep 15, 2023
+		EQ 200MG BASE	N216873	003	Sep 15, 2023

MOMETASONE FUROATE

AEROSOL, METERED; INHALATION

ASMANEX HFA

+	ORGANON LLC	0.05MG/INH	N205641	003	Aug 12, 2019
+		0.10MG/INH	N205641	001	Apr 25, 2014
+		0.20MG/INH	N205641	002	Apr 25, 2014

CREAM; TOPICAL

MOMETASONE FUROATE

AB	COSETTE	<u>0.1%</u>	<u>A077447</u>	<u>001</u>	May 22, 2006
AB	! GLENMARK PHARMS INC	<u>0.1%</u>	<u>A078541</u>	<u>001</u>	May 28, 2008
AB	TARO	<u>0.1%</u>	<u>A076679</u>	<u>001</u>	Dec 21, 2004

IMPLANT; IMPLANTATION

SINUVA

+	INTERSECT ENT INC	1.35MG	N209310	001	Dec 08, 2017
---	-------------------	--------	---------	-----	--------------

LOTION; TOPICAL

MOMETASONE FUROATE

AB	ENCUBE	<u>0.1%</u>	<u>A076499</u>	<u>001</u>	Nov 21, 2007
AB	FOUGERA PHARMS	<u>0.1%</u>	<u>A075919</u>	<u>001</u>	Nov 29, 2007
AB	GLENMARK PHARMS LTD	<u>0.1%</u>	<u>A090506</u>	<u>001</u>	Aug 09, 2010
AB	! PADAGIS ISRAEL	<u>0.1%</u>	<u>A077180</u>	<u>001</u>	Apr 06, 2005
AB	TARO	<u>0.1%</u>	<u>A076788</u>	<u>001</u>	Mar 15, 2006

OINTMENT; TOPICAL

MOMETASONE FUROATE

AB	COSETTE	<u>0.1%</u>	<u>A077401</u>	<u>001</u>	Jun 20, 2006
AB	FOUGERA PHARMS	<u>0.1%</u>	<u>A077061</u>	<u>001</u>	Mar 28, 2005
AB	! GLENMARK PHARMS INC	<u>0.1%</u>	<u>A078571</u>	<u>001</u>	May 28, 2008
AB	PADAGIS US	<u>0.1%</u>	<u>A076067</u>	<u>001</u>	Mar 18, 2002
AB	TORRENT	<u>0.1%</u>	<u>A207899</u>	<u>001</u>	Jul 13, 2018

POWDER; INHALATION

ASMANEX TWISTHALER

+	ORGANON LLC	0.11MG/INH	N021067	002	Feb 01, 2008
+		0.22MG/INH	N021067	001	Mar 30, 2005

SPRAY, METERED; NASAL

MOMETASONE FUROATE

AB	AMNEAL PHARMS	<u>0.05MG/SPRAY</u>	<u>A207989</u>	<u>001</u>	Apr 03, 2017
AB	! APOTEX	<u>0.05MG/SPRAY</u>	<u>A091161</u>	<u>001</u>	Mar 22, 2016
AB	AUROBINDO PHARMA	<u>0.05MG/SPRAY</u>	<u>A215878</u>	<u>001</u>	Mar 18, 2024

MOMETASONE FUROATE; OLOPATADINE HYDROCHLORIDE

SPRAY, METERED; NASAL

RYALTRIS

+	GLENMARK SPECLT	0.025MG/SPRAY; 0.665MG/SPRAY	N211746	001	Jan 13, 2022
---	-----------------	------------------------------	---------	-----	--------------

MONOMETHYL FUMARATE

CAPSULE, DELAYED RELEASE; ORAL

BAFIERTAM

+	BANNER LIFE SCIENCES	95MG	N210296	001	Apr 28, 2020
---	----------------------	------	---------	-----	--------------

MONTELUKAST SODIUM

GRANULE; ORAL

MONTELUKAST SODIUM

AB	AJANTA PHARMA LTD	<u>EQ 4MG BASE/PACKET</u>	<u>A203438</u>	<u>001</u>	Jul 31, 2015
AB	AUROBINDO PHARMA LTD	<u>EQ 4MG BASE/PACKET</u>	<u>A213471</u>	<u>001</u>	Feb 18, 2020
AB	DR REDDYS LABS LTD	<u>EQ 4MG BASE/PACKET</u>	<u>A202906</u>	<u>001</u>	Sep 17, 2012
AB	TORRENT	<u>EQ 4MG BASE/PACKET</u>	<u>A210431</u>	<u>001</u>	Jul 31, 2018

SINGULAIR

AB	! ORGANON	<u>EQ 4MG BASE/PACKET</u>	<u>N021409</u>	<u>001</u>	Jul 26, 2002
-----------	-----------	---------------------------	-----------------------	-------------------	--------------

TABLET; ORAL

MONTELUKAST SODIUM

AB	ACCORD HLTHCARE	<u>EQ 10MG BASE</u>	<u>A202717</u>	<u>001</u>	Sep 21, 2012
AB	AMNEAL PHARMS	<u>EQ 10MG BASE</u>	<u>A204604</u>	<u>001</u>	Sep 04, 2015
AB	ANBISON LAB	<u>EQ 10MG BASE</u>	<u>A205683</u>	<u>001</u>	Jan 12, 2016
AB	AUROBINDO PHARMA LTD	<u>EQ 10MG BASE</u>	<u>A202468</u>	<u>001</u>	Aug 03, 2012
AB	CHARTWELL MOLECULAR	<u>EQ 10MG BASE</u>	<u>A201522</u>	<u>001</u>	Aug 03, 2012
AB	CIPLA	<u>EQ 10MG BASE</u>	<u>A207463</u>	<u>001</u>	Oct 28, 2016

PRESCRIPTION DRUG PRODUCT LIST

MONTELUKAST SODIUM

TABLET; ORAL

MONTELUKAST SODIUM

<u>AB</u>	DR REDDYS LABS LTD	<u>EQ 10MG BASE</u>	<u>A201582 001</u>	Aug 06, 2012
<u>AB</u>	GLENMARK PHARMS LTD	<u>EQ 10MG BASE</u>	<u>A090926 001</u>	Aug 03, 2012
<u>AB</u>	HETERO LABS LTD V	<u>EQ 10MG BASE</u>	<u>A202843 001</u>	Sep 10, 2014
<u>AB</u>	MACLEODS PHARMS LTD	<u>EQ 10MG BASE</u>	<u>A203366 001</u>	Sep 11, 2014
<u>AB</u>	RISING	<u>EQ 10MG BASE</u>	<u>A209012 001</u>	Apr 24, 2017
<u>AB</u>	TEVA PHARMS	<u>EQ 10MG BASE</u>	<u>A078605 001</u>	Aug 03, 2012
<u>AB</u>	TORRENT PHARMS LTD	<u>EQ 10MG BASE</u>	<u>A201515 001</u>	Aug 03, 2012
<u>AB</u>	UNICHEM	<u>EQ 10MG BASE</u>	<u>A204290 001</u>	Oct 08, 2015
<u>AB</u>	UNIMARK REMEDIES LTD	<u>EQ 10MG BASE</u>	<u>A202859 001</u>	Oct 30, 2014

SINGULAIR

<u>AB</u>	<u>+</u> ORGANON	<u>EQ 10MG BASE</u>	<u>N020829 002</u>	Feb 20, 1998
-----------	------------------	---------------------	--------------------	--------------

TABLET, CHEWABLE; ORAL

MONTELUKAST SODIUM

<u>AB</u>	AMNEAL PHARMS	<u>EQ 4MG BASE</u>	<u>A205107 001</u>	Sep 04, 2020
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A205107 002</u>	Sep 04, 2020
<u>AB</u>	ANBISON LAB	<u>EQ 4MG BASE</u>	<u>A205695 001</u>	Nov 05, 2015
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A205695 002</u>	Nov 05, 2015
<u>AB</u>	AUROBINDO PHARMA LTD	<u>EQ 4MG BASE</u>	<u>A202096 001</u>	Aug 03, 2012
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A202096 002</u>	Aug 03, 2012
<u>AB</u>	CHARTWELL MOLECULAR	<u>EQ 4MG BASE</u>	<u>A207464 001</u>	Dec 06, 2018
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A207464 002</u>	Dec 06, 2018
<u>AB</u>	DR REDDYS LABS LTD	<u>EQ 4MG BASE</u>	<u>A201581 001</u>	Aug 06, 2012
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A201581 002</u>	Aug 06, 2012
<u>AB</u>	HETERO LABS LTD V	<u>EQ 4MG BASE</u>	<u>A204093 001</u>	May 22, 2015
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A204093 002</u>	May 22, 2015
<u>AB</u>	LANNETT CO INC	<u>EQ 4MG BASE</u>	<u>A200405 001</u>	Aug 03, 2012
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A200405 002</u>	Aug 03, 2012
<u>AB</u>	MACLEODS PHARMS LTD	<u>EQ 4MG BASE</u>	<u>A203582 001</u>	Mar 12, 2015
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A203582 002</u>	Mar 12, 2015
<u>AB</u>	RISING	<u>EQ 4MG BASE</u>	<u>A209011 001</u>	Apr 18, 2017
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A209011 002</u>	Apr 18, 2017
<u>AB</u>	TEVA PHARMS	<u>EQ 4MG BASE</u>	<u>A078723 001</u>	Aug 03, 2012
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A078723 002</u>	Aug 03, 2012
<u>AB</u>	TORRENT PHARMS LTD	<u>EQ 4MG BASE</u>	<u>A090984 001</u>	Aug 03, 2012
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A090984 002</u>	Aug 03, 2012
<u>AB</u>	UNIMARK REMEDIES LTD	<u>EQ 4MG BASE</u>	<u>A203037 001</u>	Oct 30, 2014
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A203037 002</u>	Oct 30, 2014

SINGULAIR

<u>AB</u>	<u>+</u> ORGANON	<u>EQ 4MG BASE</u>	<u>N020830 002</u>	Mar 03, 2000
<u>AB</u>	<u>+</u> !	<u>EQ 5MG BASE</u>	<u>N020830 001</u>	Feb 20, 1998

MORPHINE SULFATE

CAPSULE, EXTENDED RELEASE; ORAL

MORPHINE SULFATE

	ACTAVIS ELIZABETH	30MG	A079040 001	Jan 16, 2013
		45MG	A079040 002	Jan 16, 2013
		60MG	A079040 003	Jan 16, 2013
		75MG	A079040 004	Jan 16, 2013
		90MG	A079040 005	Jan 16, 2013
!		120MG	A079040 006	Jan 16, 2013
	UPSHER SMITH LABS	10MG	A202104 001	Jun 03, 2013
		20MG	A202104 002	Jun 03, 2013
		30MG	A202104 003	Jun 03, 2013
		50MG	A202104 004	Jun 03, 2013
		60MG	A202104 005	Jun 03, 2013
		80MG	A202104 006	Jun 03, 2013
!		100MG	A202104 007	Jun 03, 2013

INJECTABLE; INJECTION

DURAMORPH PF

<u>AP</u>	<u>+</u> ! HIKMA	<u>0.5MG/ML</u>	<u>N018565 001</u>	Sep 18, 1984
<u>AP</u>	<u>+</u> !	<u>1MG/ML</u>	<u>N018565 002</u>	Sep 18, 1984

INFUMORPH

<u>AP</u>	<u>+</u> ! HIKMA	<u>10MG/ML</u>	<u>N018565 003</u>	Jul 19, 1991
<u>AP</u>	<u>+</u> !	<u>25MG/ML</u>	<u>N018565 004</u>	Jul 19, 1991

MITIGO

<u>AP</u>	PIRAMAL CRITICAL	<u>10MG/ML</u>	<u>A204393 001</u>	Jul 16, 2018
<u>AP</u>		<u>25MG/ML</u>	<u>A204393 002</u>	Jul 16, 2018

PRESCRIPTION DRUG PRODUCT LIST

MORPHINE SULFATE

INJECTABLE; INJECTION

MORPHINE SULFATE

<u>AP</u>	HIKMA	<u>2MG/ML</u>	<u>A211452 001</u>	Jan 12, 2023
<u>AP</u>	!	<u>4MG/ML</u>	<u>A205758 001</u>	May 21, 2015
<u>AP</u>		<u>4MG/ML</u>	<u>A211452 002</u>	Jan 12, 2023
<u>AP</u>	!	<u>8MG/ML</u>	<u>A205758 002</u>	May 21, 2015
<u>AP</u>		<u>8MG/ML</u>	<u>A211452 003</u>	Jan 12, 2023
<u>AP</u>		<u>10MG/ML</u>	<u>A205758 003</u>	May 21, 2015
<u>AP</u>		<u>10MG/ML</u>	<u>A211452 004</u>	Jan 12, 2023
<u>AP</u>	HOSPIRA	<u>0.5MG/ML</u>	<u>A073509 001</u>	Sep 30, 1992
<u>AP</u>		<u>1MG/ML</u>	<u>A073510 001</u>	Sep 30, 1992
<u>AP</u>	+!	<u>2MG/ML</u>	<u>N202515 001</u>	Nov 14, 2011
<u>AP</u>	+	<u>4MG/ML</u>	<u>N202515 002</u>	Nov 14, 2011
<u>AP</u>	+	<u>8MG/ML</u>	<u>N202515 003</u>	Nov 14, 2011
<u>AP</u>	+	<u>10MG/ML</u>	<u>N202515 004</u>	Nov 14, 2011
	HIKMA	15MG/ML	A211452 005	Jan 12, 2023
	+!	HOSPIRA INC	N202515 006	Apr 29, 2021
		INTL MEDICATION SYS	A202861 001	Apr 29, 2021

SOLUTION; INTRAMUSCULAR, INTRAVENOUS

MORPHINE SULFATE

	+!	FRESENIUS KABI USA	2MG/ML (2MG/ML)	N204223 001	Oct 30, 2013
	+!		4MG/ML (4MG/ML)	N204223 002	Oct 30, 2013
	+!		10MG/ML (10MG/ML)	N204223 005	Oct 30, 2013

SOLUTION; ORAL

MORPHINE SULFATE

<u>AA</u>	+	HIKMA	<u>10MG/5ML</u>	<u>N022195 001</u>	Mar 17, 2008
<u>AA</u>	+		<u>20MG/5ML</u>	<u>N022195 002</u>	Mar 17, 2008
<u>AA</u>	+!		<u>100MG/5ML</u>	<u>N022195 003</u>	Jan 25, 2010
<u>AA</u>		PADAGIS US	<u>100MG/5ML</u>	<u>A201574 001</u>	Aug 06, 2012
<u>AA</u>		RHODES PHARMS	<u>10MG/5ML</u>	<u>A206308 001</u>	Jun 22, 2017
<u>AA</u>			<u>20MG/5ML</u>	<u>A206420 001</u>	Jul 12, 2016
<u>AA</u>			<u>100MG/5ML</u>	<u>A206308 002</u>	Jun 22, 2017
<u>AA</u>		SPECGX LLC	<u>100MG/5ML</u>	<u>A202348 001</u>	Jul 15, 2011
<u>AA</u>		TRIS PHARMA INC	<u>10MG/5ML</u>	<u>A203518 001</u>	May 12, 2015
<u>AA</u>			<u>100MG/5ML</u>	<u>A203518 002</u>	May 12, 2015
<u>AA</u>		WINDER LABS LLC	<u>10MG/5ML</u>	<u>A211454 001</u>	Feb 12, 2021
<u>AA</u>			<u>20MG/5ML</u>	<u>A211454 002</u>	Feb 12, 2021
<u>AA</u>			<u>100MG/5ML</u>	<u>A211454 003</u>	Feb 12, 2021

TABLET; ORAL

MORPHINE SULFATE

<u>AB</u>		ALKEM LABS LTD	<u>15MG</u>	<u>A212451 001</u>	Dec 03, 2020
<u>AB</u>			<u>30MG</u>	<u>A212451 002</u>	Dec 03, 2020
<u>AB</u>	+	HIKMA	<u>15MG</u>	<u>N022207 001</u>	Mar 17, 2008
<u>AB</u>	+!		<u>30MG</u>	<u>N022207 002</u>	Mar 17, 2008
<u>AB</u>		SPECGX LLC	<u>15MG</u>	<u>A215194 001</u>	Aug 21, 2023
<u>AB</u>			<u>30MG</u>	<u>A215194 002</u>	Aug 21, 2023
<u>AB</u>		UPSHER SMITH LABS	<u>15MG</u>	<u>A210610 001</u>	Jul 22, 2019
<u>AB</u>			<u>30MG</u>	<u>A210610 002</u>	Jul 22, 2019

TABLET, EXTENDED RELEASE; ORAL

MORPHINE SULFATE

<u>AB</u>		ACTAVIS ELIZABETH	<u>15MG</u>	<u>A203849 001</u>	Apr 06, 2015
<u>AB</u>			<u>30MG</u>	<u>A203849 002</u>	Apr 06, 2015
<u>AB</u>			<u>60MG</u>	<u>A203849 003</u>	Apr 06, 2015
<u>AB</u>			<u>100MG</u>	<u>A203849 004</u>	Apr 06, 2015
<u>AB</u>			<u>200MG</u>	<u>A203849 005</u>	Apr 06, 2015
<u>AB</u>		DAVA PHARMS INC	<u>15MG</u>	<u>A075407 001</u>	Jan 28, 2000
<u>AB</u>		NOVEL LABS INC	<u>15MG</u>	<u>A203602 001</u>	Dec 16, 2015
<u>AB</u>			<u>30MG</u>	<u>A203602 002</u>	Dec 16, 2015
<u>AB</u>			<u>60MG</u>	<u>A203602 003</u>	Dec 16, 2015
<u>AB</u>			<u>100MG</u>	<u>A203602 004</u>	Dec 16, 2015
<u>AB</u>			<u>200MG</u>	<u>A203602 005</u>	Dec 16, 2015
<u>AB</u>		RHODES PHARMS	<u>15MG</u>	<u>A074862 001</u>	Jul 07, 1998
<u>AB</u>			<u>30MG</u>	<u>A074862 002</u>	Jul 07, 1998
<u>AB</u>			<u>60MG</u>	<u>A074862 003</u>	Jul 07, 1998
<u>AB</u>			<u>100MG</u>	<u>A074769 001</u>	Jul 02, 1998
<u>AB</u>			<u>200MG</u>	<u>A074769 002</u>	Jul 02, 1998
<u>AB</u>		SPECGX LLC	<u>15MG</u>	<u>A076412 001</u>	Jul 31, 2003
<u>AB</u>			<u>30MG</u>	<u>A076412 002</u>	Jul 31, 2003
<u>AB</u>			<u>60MG</u>	<u>A076412 003</u>	Jul 31, 2003
<u>AB</u>			<u>100MG</u>	<u>A076438 001</u>	Jul 03, 2003
<u>AB</u>			<u>200MG</u>	<u>A076438 002</u>	Jul 03, 2003

PRESCRIPTION DRUG PRODUCT LIST

MORPHINE SULFATE

TABLET, EXTENDED RELEASE;ORAL

MORPHINE SULFATE

<u>AB</u>	STRIDES PHARMA	<u>15MG</u>	<u>A075295 001</u>	Oct 28, 1998
<u>AB</u>		<u>30MG</u>	<u>A075295 002</u>	Oct 28, 1998
<u>AB</u>		<u>60MG</u>	<u>A075295 003</u>	Oct 28, 1998
<u>AB</u>		<u>100MG</u>	<u>A075295 004</u>	Sep 15, 2000
<u>AB</u>		<u>200MG</u>	<u>A075295 005</u>	Sep 15, 2000
<u>AB</u>	SUN PHARM INDS LTD	<u>15MG</u>	<u>A078761 001</u>	May 11, 2012
<u>AB</u>		<u>30MG</u>	<u>A078761 002</u>	May 11, 2012
<u>AB</u>		<u>60MG</u>	<u>A078761 003</u>	May 11, 2012
<u>AB</u>		<u>100MG</u>	<u>A078761 004</u>	May 11, 2012
<u>AB</u>		<u>200MG</u>	<u>A078761 005</u>	May 11, 2012

MS CONTIN

<u>AB</u>	+ PURDUE PHARMA LP	<u>15MG</u>	<u>N019516 003</u>	Sep 12, 1989
<u>AB</u>	+	<u>30MG</u>	<u>N019516 001</u>	May 29, 1987
<u>AB</u>	+	<u>60MG</u>	<u>N019516 002</u>	Apr 08, 1988
<u>AB</u>	+!	<u>100MG</u>	<u>N019516 004</u>	Jan 16, 1990
<u>AB</u>	+	<u>200MG</u>	<u>N019516 005</u>	Nov 08, 1993

MOTIXAFORTIDE ACETATE

POWDER;SUBCUTANEOUS

APHEXDA

+! BIOLINERX LTD EQ 62MG BASE/VIAL N217159 001 Sep 08, 2023

MOXIDECTIN

TABLET;ORAL

MOXIDECTIN

+! MDGH 2MG N210867 001 Jun 13, 2018

MOXIFLOXACIN HYDROCHLORIDE

SOLUTION;INTRAVENOUS

MOXIFLOXACIN HYDROCHLORIDE

+! FRESENIUS KABI USA EQ 400MG BASE/250ML (EQ 1.6MG BASE/ML) N205572 001 Apr 03, 2015

MOXIFLOXACIN HYDROCHLORIDE IN SODIUM CHLORIDE 0.8% IN PLASTIC CONTAINER

! MYLAN LABS LTD 400MG/250ML (1.6MG/ML) A205833 001 May 05, 2017

SOLUTION/DROPS;OPHTHALMIC

MOXIFLOXACIN HYDROCHLORIDE

<u>AT1</u>	ADAPTIS	<u>EQ 0.5% BASE</u>	<u>A202525 001</u>	Mar 06, 2015
<u>AT1</u>	ALEMBIC	<u>EQ 0.5% BASE</u>	<u>A209469 001</u>	Feb 13, 2019
<u>AT1</u>	APOTEX	<u>EQ 0.5% BASE</u>	<u>A090080 001</u>	Jun 30, 2017
<u>AT1</u>	EUGIA PHARMA	<u>EQ 0.5% BASE</u>	<u>A206242 001</u>	Oct 04, 2017
<u>AT1</u>	GLAND PHARMA LTD	<u>EQ 0.5% BASE</u>	<u>A208778 001</u>	Mar 30, 2020
<u>AT1</u>	LUPIN LTD	<u>EQ 0.5% BASE</u>	<u>A202867 001</u>	Sep 04, 2014
<u>AT1</u>	SOMERSET THERAPS LLC	<u>EQ 0.5% BASE</u>	<u>A209698 001</u>	Nov 14, 2024
<u>AT1</u>	UPSHER SMITH LABS	<u>EQ 0.5% BASE</u>	<u>A212616 001</u>	Feb 10, 2021

VIGAMOXAT1 +! HARROW EYE EQ 0.5% BASE N021598 001 Apr 15, 2003

MOXIFLOXACIN HYDROCHLORIDE

! LUPIN LTD EQ 0.5% BASE A204079 001 May 28, 2015

TABLET;ORAL

MOXIFLOXACIN HYDROCHLORIDE

<u>AB</u>	AUROBINDO PHARMA LTD	<u>EQ 400MG BASE</u>	<u>A202632 001</u>	Mar 04, 2014
<u>AB</u>	CHARTWELL RX	<u>EQ 400MG BASE</u>	<u>A207285 001</u>	Feb 13, 2017
<u>AB</u>	CROSSMEDIKA SA	<u>EQ 400MG BASE</u>	<u>A205348 001</u>	Jan 14, 2016
<u>AB</u>	! DR REDDYS	<u>EQ 400MG BASE</u>	<u>A076938 001</u>	Mar 04, 2014
<u>AB</u>	HETERO LABS LTD V	<u>EQ 400MG BASE</u>	<u>A204836 001</u>	Mar 02, 2023
<u>AB</u>	MSN	<u>EQ 400MG BASE</u>	<u>A208682 001</u>	Sep 22, 2017
<u>AB</u>	TEVA PHARMS USA	<u>EQ 400MG BASE</u>	<u>A077437 001</u>	Feb 18, 2014

MUPIROICIN

OINTMENT;TOPICAL

MUPIROICIN

<u>AB</u>	FOUGERA PHARMS	<u>2%</u>	<u>A065192 001</u>	Nov 30, 2005
<u>AB</u>	GLENMARK PHARMS	<u>2%</u>	<u>A090480 001</u>	Jun 08, 2011
<u>AB</u>	! PADAGIS ISRAEL	<u>2%</u>	<u>A065123 001</u>	Nov 07, 2003
<u>AB</u>	TARO	<u>2%</u>	<u>A065170 001</u>	Sep 23, 2005
<u>AB</u>	TEVA	<u>2%</u>	<u>A065085 001</u>	Nov 07, 2003

CENTANY

BX PADAGIS US 2% N050788 001 Dec 04, 2002

PRESCRIPTION DRUG PRODUCT LIST

MUPIROCIN CALCIUM

CREAM; TOPICAL

MUPIROCIN

<u>AB</u>	ALEMBIC	<u>EQ 2% BASE</u>	<u>A213053</u>	<u>001</u>	Nov 16, 2021
<u>AB</u>	AMNEAL	<u>EQ 2% BASE</u>	<u>A214811</u>	<u>001</u>	Nov 15, 2022
<u>AB</u>	ENCUBE	<u>EQ 2% BASE</u>	<u>A213076</u>	<u>001</u>	Aug 31, 2021
<u>AB</u>	! GLENMARK PHARMS INC	<u>EQ 2% BASE</u>	<u>A201587</u>	<u>001</u>	Jan 24, 2013
<u>AB</u>	TARO	<u>EQ 2% BASE</u>	<u>A207116</u>	<u>001</u>	Apr 27, 2020

MYCOPHENOLATE MOFETIL

CAPSULE; ORAL

CELLCEPT

<u>AB</u>	+! ROCHE PALO	<u>250MG</u>	<u>N050722</u>	<u>001</u>	May 03, 1995
-----------	---------------	--------------	----------------	------------	--------------

MYCOPHENOLATE MOFETIL

<u>AB</u>	ACCORD HLTHCARE	<u>250MG</u>	<u>A090253</u>	<u>001</u>	May 04, 2009
<u>AB</u>	ALKEM LABS LTD	<u>250MG</u>	<u>A200197</u>	<u>001</u>	Jun 13, 2013
<u>AB</u>	AUROBINDO PHARMA LTD	<u>250MG</u>	<u>A217828</u>	<u>001</u>	Jan 05, 2024
<u>AB</u>	CONCORD BIOTECH LTD	<u>250MG</u>	<u>A210181</u>	<u>001</u>	Jan 08, 2019
<u>AB</u>	HETERO LABS LTD V	<u>250MG</u>	<u>A207022</u>	<u>001</u>	Oct 22, 2024
<u>AB</u>	HIKMA	<u>250MG</u>	<u>A065410</u>	<u>001</u>	Jul 29, 2008
<u>AB</u>	MYLAN	<u>250MG</u>	<u>A065520</u>	<u>001</u>	May 04, 2009
<u>AB</u>	SANDOZ	<u>250MG</u>	<u>A065379</u>	<u>001</u>	Oct 15, 2008
<u>AB</u>	STRIDES PHARMA	<u>250MG</u>	<u>A090055</u>	<u>001</u>	Jun 10, 2010
<u>AB</u>	TEVA PHARMS	<u>250MG</u>	<u>A065491</u>	<u>001</u>	May 06, 2009
<u>AB</u>	WUXI	<u>250MG</u>	<u>A214079</u>	<u>001</u>	Nov 07, 2024
<u>AB</u>	ZHEJIANG HISUN PHARM	<u>250MG</u>	<u>A204077</u>	<u>001</u>	Nov 13, 2017

FOR SUSPENSION; ORAL

CELLCEPT

<u>AB</u>	+! ROCHE PALO	<u>200MG/ML</u>	<u>N050759</u>	<u>001</u>	Oct 01, 1998
-----------	---------------	-----------------	----------------	------------	--------------

MYCOPHENOLATE MOFETIL

<u>AB</u>	ALKEM LABS LTD	<u>200MG/ML</u>	<u>A203005</u>	<u>001</u>	Nov 14, 2014
<u>AB</u>	AMNEAL	<u>200MG/ML</u>	<u>A214871</u>	<u>001</u>	Nov 02, 2021
<u>AB</u>	AUROBINDO PHARMA LTD	<u>200MG/ML</u>	<u>A218227</u>	<u>001</u>	Sep 26, 2024
<u>AB</u>	LANNETT CO INC	<u>200MG/ML</u>	<u>A214525</u>	<u>001</u>	Jul 29, 2021
<u>AB</u>	RISING	<u>200MG/ML</u>	<u>A213955</u>	<u>001</u>	Sep 11, 2023
<u>AB</u>	STRIDES PHARMA	<u>200MG/ML</u>	<u>A212634</u>	<u>001</u>	Aug 29, 2023
<u>AB</u>	TEVA PHARMS USA	<u>200MG/ML</u>	<u>A211272</u>	<u>001</u>	Jan 25, 2022
<u>AB</u>	VISTAPHARM	<u>200MG/ML</u>	<u>A210370</u>	<u>001</u>	Feb 12, 2019

SUSPENSION; ORAL

MYHIBBIN

	+! AZURITY	200MG/ML	N216482	001	May 01, 2024
--	------------	----------	---------	-----	--------------

TABLET; ORAL

CELLCEPT

<u>AB</u>	+! ROCHE PALO	<u>500MG</u>	<u>N050723</u>	<u>001</u>	Jun 19, 1997
-----------	---------------	--------------	----------------	------------	--------------

MYCOPHENOLATE MOFETIL

<u>AB</u>	ACCORD HLTHCARE	<u>500MG</u>	<u>A065416</u>	<u>001</u>	May 04, 2009
<u>AB</u>	ALKEM LABS LTD	<u>500MG</u>	<u>A091249</u>	<u>001</u>	Nov 04, 2011
<u>AB</u>	AUROBINDO PHARMA	<u>500MG</u>	<u>A217937</u>	<u>001</u>	Feb 20, 2024
<u>AB</u>	CONCORD BIOTECH LTD	<u>500MG</u>	<u>A212087</u>	<u>001</u>	Jul 31, 2020
<u>AB</u>	HETERO LABS LTD V	<u>500MG</u>	<u>A208119</u>	<u>001</u>	Oct 22, 2024
<u>AB</u>	HIKMA	<u>500MG</u>	<u>A065413</u>	<u>001</u>	Jul 29, 2008
<u>AB</u>	MYLAN	<u>500MG</u>	<u>A065521</u>	<u>001</u>	May 04, 2009
<u>AB</u>	SANDOZ	<u>500MG</u>	<u>A065451</u>	<u>001</u>	Oct 15, 2008
<u>AB</u>	STRIDES PHARMA	<u>500MG</u>	<u>A090456</u>	<u>001</u>	Jun 10, 2010
<u>AB</u>	ZHEJIANG HISUN PHARM	<u>500MG</u>	<u>A204076</u>	<u>001</u>	Nov 16, 2017

MYCOPHENOLATE MOFETIL HYDROCHLORIDE

INJECTABLE; INJECTION

CELLCEPT

<u>AP</u>	+! ROCHE PALO	<u>500MG/VIAL</u>	<u>N050758</u>	<u>001</u>	Aug 12, 1998
-----------	---------------	-------------------	----------------	------------	--------------

MYCOPHENOLATE MOFETIL HYDROCHLORIDE

<u>AP</u>	BPI LABS	<u>500MG/VIAL</u>	<u>A214283</u>	<u>001</u>	Jun 01, 2023
<u>AP</u>	ENDO OPERATIONS	<u>500MG/VIAL</u>	<u>A203575</u>	<u>001</u>	Oct 28, 2016
<u>AP</u>	MEITHEAL	<u>500MG/VIAL</u>	<u>A212130</u>	<u>001</u>	Jan 15, 2021
<u>AP</u>	MYLAN LABS LTD	<u>500MG/VIAL</u>	<u>A203859</u>	<u>001</u>	Mar 31, 2017
<u>AP</u>	RISING	<u>500MG/VIAL</u>	<u>A204043</u>	<u>001</u>	Feb 28, 2017
<u>AP</u>	STERISCIENCE	<u>500MG/VIAL</u>	<u>A216390</u>	<u>001</u>	Dec 23, 2022
<u>AP</u>	SPECLTS				
<u>AP</u>	ZYDUS PHARMS	<u>500MG/VIAL</u>	<u>A204473</u>	<u>001</u>	Aug 31, 2017

PRESCRIPTION DRUG PRODUCT LIST

MYCOPHENOLIC SODIUM

TABLET, DELAYED RELEASE;ORAL

MYCOPHENOLIC SODIUM

<u>AB</u>	ACCORD HLTHCARE	<u>EQ 180MG BASE</u>	<u>A202555 001</u>	Aug 23, 2017
<u>AB</u>		<u>EQ 360MG BASE</u>	<u>A202555 002</u>	Aug 23, 2017
<u>AB</u>	ALKEM LABS LTD	<u>EQ 180MG BASE</u>	<u>A208315 001</u>	Sep 23, 2021
<u>AB</u>		<u>EQ 360MG BASE</u>	<u>A208315 002</u>	Sep 23, 2021
<u>AB</u>	AMTA	<u>EQ 180MG BASE</u>	<u>A214376 001</u>	Feb 10, 2021
<u>AB</u>		<u>EQ 360MG BASE</u>	<u>A214376 002</u>	Feb 10, 2021
<u>AB</u>	APOTEX INC	<u>EQ 180MG BASE</u>	<u>A091558 001</u>	Aug 21, 2012
<u>AB</u>		<u>EQ 360MG BASE</u>	<u>A091558 002</u>	Aug 19, 2014
<u>AB</u>	AUROBINDO PHARMA LTD	<u>EQ 180MG BASE</u>	<u>A218603 001</u>	Feb 27, 2024
<u>AB</u>		<u>EQ 360MG BASE</u>	<u>A218603 002</u>	Feb 27, 2024
<u>AB</u>	BIOCON PHARMA	<u>EQ 180MG BASE</u>	<u>A214630 001</u>	Nov 29, 2021
<u>AB</u>		<u>EQ 180MG BASE</u>	<u>A217031 001</u>	Nov 29, 2023
<u>AB</u>		<u>EQ 360MG BASE</u>	<u>A214630 002</u>	Nov 29, 2021
<u>AB</u>		<u>EQ 360MG BASE</u>	<u>A217031 002</u>	Nov 29, 2023
<u>AB</u>	CONCORD BIOTECH LTD	<u>EQ 180MG BASE</u>	<u>A211173 001</u>	Dec 13, 2019
<u>AB</u>		<u>EQ 360MG BASE</u>	<u>A211173 002</u>	Dec 13, 2019
<u>AB</u>	RK PHARMA	<u>EQ 180MG BASE</u>	<u>A091248 002</u>	Jan 08, 2014
<u>AB</u>		<u>EQ 360MG BASE</u>	<u>A091248 001</u>	Jan 08, 2014
<u>AB</u>	TWI PHARMS	<u>EQ 180MG BASE</u>	<u>A214289 001</u>	Nov 03, 2021
<u>AB</u>		<u>EQ 360MG BASE</u>	<u>A214289 002</u>	Nov 03, 2021
<u>AB</u>	WANBANG BIOPHARMS	<u>EQ 180MG BASE</u>	<u>A216637 001</u>	May 29, 2024
<u>AB</u>		<u>EQ 360MG BASE</u>	<u>A216637 002</u>	May 29, 2024
<u>MYFORTIC</u>				
<u>AB</u>	+ NOVARTIS	<u>EQ 180MG BASE</u>	<u>N050791 001</u>	Feb 27, 2004
<u>AB</u>	+!	<u>EQ 360MG BASE</u>	<u>N050791 002</u>	Feb 27, 2004

NABUMETONE

TABLET;ORAL

NABUMETONE

<u>AB</u>	ANNORA PHARMA	<u>500MG</u>	<u>A090445 001</u>	Jan 12, 2011
<u>AB</u>		<u>750MG</u>	<u>A090445 002</u>	Jan 12, 2011
<u>AB</u>	CHARTWELL MOLECULES	<u>500MG</u>	<u>A076009 001</u>	Jan 24, 2003
<u>AB</u>		<u>750MG</u>	<u>A076009 002</u>	Jan 24, 2003
<u>AB</u>	CHARTWELL RX	<u>500MG</u>	<u>A075280 001</u>	Feb 25, 2002
<u>AB</u>		<u>750MG</u>	<u>A075280 002</u>	Feb 25, 2002
<u>AB</u>	INVAGEN PHARMS	<u>500MG</u>	<u>A078671 001</u>	Mar 07, 2008
<u>AB</u>	!	<u>750MG</u>	<u>A078671 002</u>	Mar 07, 2008
<u>AB</u>	LGM PHARMA	<u>500MG</u>	<u>A203166 001</u>	Aug 30, 2019
<u>AB</u>		<u>750MG</u>	<u>A203166 002</u>	Aug 30, 2019
<u>AB</u>	NOVITIUM PHARMA	<u>500MG</u>	<u>A219118 001</u>	Jan 07, 2025
<u>AB</u>		<u>750MG</u>	<u>A219118 002</u>	Jan 07, 2025
<u>AB</u>	SCIEGEN PHARMS INC	<u>500MG</u>	<u>A078420 001</u>	Sep 24, 2008
<u>AB</u>		<u>750MG</u>	<u>A078420 002</u>	Sep 24, 2008
<u>AB</u>	WATSON LABS	<u>500MG</u>	<u>A091083 001</u>	Jun 13, 2011
<u>AB</u>		<u>750MG</u>	<u>A091083 002</u>	Jun 13, 2011
<u>AB</u>	LGM PHARMA	1GM	A203166 003	Aug 30, 2019

NADOLOL

TABLET;ORAL

NADOLOL

<u>AB</u>	ALEMBIC	<u>20MG</u>	<u>A211763 001</u>	Jun 02, 2023
<u>AB</u>		<u>40MG</u>	<u>A211763 002</u>	Jun 02, 2023
<u>AB</u>		<u>80MG</u>	<u>A211763 003</u>	Jun 02, 2023
<u>AB</u>	AMNEAL PHARMS CO	<u>20MG</u>	<u>A208832 001</u>	Jun 02, 2017
<u>AB</u>		<u>40MG</u>	<u>A208832 002</u>	Jun 02, 2017
<u>AB</u>		<u>80MG</u>	<u>A208832 003</u>	Jun 02, 2017
<u>AB</u>	AUROBINDO PHARMA	<u>20MG</u>	<u>A201893 003</u>	Jun 07, 2022
<u>AB</u>		<u>40MG</u>	<u>A201893 001</u>	Sep 16, 2015
<u>AB</u>		<u>80MG</u>	<u>A201893 002</u>	Sep 16, 2015
<u>AB</u>	BEXIMCO PHARMS USA	<u>20MG</u>	<u>A210955 001</u>	Jul 23, 2018
<u>AB</u>		<u>40MG</u>	<u>A210955 002</u>	Jul 23, 2018
<u>AB</u>		<u>80MG</u>	<u>A210955 003</u>	Jul 23, 2018
<u>AB</u>	CHARTWELL RX	<u>20MG</u>	<u>A209309 001</u>	Oct 05, 2017
<u>AB</u>		<u>40MG</u>	<u>A209309 002</u>	Oct 05, 2017
<u>AB</u>		<u>80MG</u>	<u>A209309 003</u>	Oct 05, 2017
<u>AB</u>	INVAGEN PHARMS	<u>20MG</u>	<u>A203455 001</u>	Dec 18, 2015
<u>AB</u>		<u>40MG</u>	<u>A203455 002</u>	Dec 18, 2015
<u>AB</u>	!	<u>80MG</u>	<u>A203455 003</u>	Dec 18, 2015
<u>AB</u>	RISING	<u>20MG</u>	<u>A074172 001</u>	Oct 31, 1993
<u>AB</u>		<u>40MG</u>	<u>A074172 002</u>	Oct 31, 1993

PRESCRIPTION DRUG PRODUCT LIST

NADOLOL

TABLET; ORAL

NADOLOL

<u>AB</u>		<u>80MG</u>	<u>A074172 003</u>	Oct 31, 1993
<u>AB</u>	RK PHARMA	<u>20MG</u>	<u>A212856 001</u>	Sep 13, 2019
<u>AB</u>		<u>40MG</u>	<u>A212856 002</u>	Sep 13, 2019
<u>AB</u>		<u>80MG</u>	<u>A212856 003</u>	Sep 13, 2019
<u>AB</u>	SANDOZ	<u>20MG</u>	<u>A074501 001</u>	Nov 09, 1995
<u>AB</u>		<u>40MG</u>	<u>A074501 002</u>	Nov 09, 1995
<u>AB</u>		<u>80MG</u>	<u>A074501 003</u>	Nov 09, 1995
<u>AB</u>	ZYDUS PHARMS	<u>20MG</u>	<u>A207761 001</u>	Jul 28, 2017
<u>AB</u>		<u>40MG</u>	<u>A207761 002</u>	Jul 28, 2017
<u>AB</u>		<u>80MG</u>	<u>A207761 003</u>	Jul 28, 2017

NAFARELIN ACETATE

SPRAY, METERED; NASAL

SYNAREL

+! PFIZER

EQ 0.2MG BASE/SPRAY

N019886 001 Feb 13, 1990

NAFCILLIN SODIUM

INJECTABLE; INJECTION

NAFCILLIN SODIUM

<u>AP</u>	ANTIBIOTICE	<u>EQ 1GM BASE/VIAL</u>	<u>A090560 001</u>	Oct 03, 2011
<u>AP</u>		<u>EQ 2GM BASE/VIAL</u>	<u>A090560 002</u>	Oct 03, 2011
<u>AP</u>	! EUGIA PHARMA SPECLTS	<u>EQ 1GM BASE/VIAL</u>	<u>A091613 001</u>	Dec 26, 2012
<u>AP</u>	!	<u>EQ 2GM BASE/VIAL</u>	<u>A091613 002</u>	Dec 26, 2012
<u>AP</u>	!	<u>EQ 10GM BASE/VIAL</u>	<u>A091614 001</u>	Dec 26, 2012
<u>AP</u>	FRESENIUS	<u>EQ 10GM BASE/VIAL</u>	<u>A206761 001</u>	Jun 02, 2020
<u>AP</u>	ISTITUTO BIO ITA SPA	<u>EQ 1GM BASE/VIAL</u>	<u>A090002 001</u>	Jun 30, 2011
<u>AP</u>		<u>EQ 2GM BASE/VIAL</u>	<u>A090002 002</u>	Jun 30, 2011
<u>AP</u>		<u>EQ 10GM BASE/VIAL</u>	<u>A090005 001</u>	Apr 20, 2011
<u>AP</u>	SAGENT PHARMS	<u>EQ 1GM BASE/VIAL</u>	<u>A090582 001</u>	Aug 24, 2012
<u>AP</u>		<u>EQ 2GM BASE/VIAL</u>	<u>A090582 002</u>	Aug 24, 2012
<u>AP</u>		<u>EQ 10GM BASE/VIAL</u>	<u>A090580 001</u>	Aug 24, 2012
<u>AP</u>	STERISCIENCE SPECLTS	<u>EQ 1GM BASE/VIAL</u>	<u>A200002 001</u>	Apr 07, 2014
<u>AP</u>		<u>EQ 2GM BASE/VIAL</u>	<u>A200002 002</u>	Apr 07, 2014
	NALLPEN IN PLASTIC CONTAINER			
	+! BAXTER HLTHCARE	EQ 20MG BASE/ML	N050655 001	Oct 31, 1989
	+!	EQ 2GM BASE/100ML	N050655 002	Oct 31, 1989

NAFTIFINE HYDROCHLORIDE

CREAM; TOPICAL

NAFTIFINE HYDROCHLORIDE

<u>AB</u>	AMNEAL	<u>2%</u>	<u>A206960 001</u>	Apr 10, 2017
<u>AB</u>	TARO	<u>2%</u>	<u>A206901 001</u>	Jan 06, 2016
<u>AB</u>	XIROMED	<u>2%</u>	<u>A210038 001</u>	Sep 22, 2020

NAFTIN

<u>AB</u>	+! LEGACY PHARMA	<u>2%</u>	<u>N019599 002</u>	Jan 13, 2012
	NAFTIFINE HYDROCHLORIDE			
	! TARO	1%	A205975 001	Sep 08, 2016

GEL; TOPICAL

NAFTIFINE HYDROCHLORIDE

<u>AB</u>	AMNEAL	<u>1%</u>	<u>A206165 001</u>	Mar 20, 2019
<u>AB</u>	TARO	<u>2%</u>	<u>A208201 001</u>	Apr 10, 2019

NAFTIN

<u>AB</u>	+! LEGACY PHARMA	<u>1%</u>	<u>N019356 001</u>	Jun 18, 1990
<u>AB</u>	+!	<u>2%</u>	<u>N204286 001</u>	Jun 27, 2013

NALBUPHINE HYDROCHLORIDE

INJECTABLE; INJECTION

NALBUPHINE HYDROCHLORIDE

<u>AP</u>	! HOSPIRA	<u>10MG/ML</u>	<u>A070914 001</u>	Feb 03, 1989
<u>AP</u>	!	<u>10MG/ML</u>	<u>A070915 001</u>	Feb 03, 1989
<u>AP</u>	!	<u>20MG/ML</u>	<u>A070916 001</u>	Feb 03, 1989
<u>AP</u>	!	<u>20MG/ML</u>	<u>A070918 001</u>	Feb 03, 1989
<u>AP</u>	SOMERSET THERAPS LLC	<u>10MG/ML</u>	<u>A216049 001</u>	Sep 19, 2024
<u>AP</u>		<u>10MG/ML</u>	<u>A216050 001</u>	Sep 19, 2024
<u>AP</u>		<u>20MG/ML</u>	<u>A216049 002</u>	Sep 19, 2024
<u>AP</u>		<u>20MG/ML</u>	<u>A216050 002</u>	Sep 19, 2024

PRESCRIPTION DRUG PRODUCT LIST

NALDEMEDINE TOSYLATE

TABLET;ORAL

SYMPROIC

+! BDSI EQ 0.2MG BASE N208854 001 Mar 23, 2017

NALMEFENE HYDROCHLORIDE

SOLUTION;INTRAMUSCULAR, INTRAVENOUS, SUBCUTANEOUS

NALMEFENE HYDROCHLORIDEAP CHENGDU SHUODE EQ 2MG BASE/2ML (EQ 1MG BASE/ML) A216007 002 Nov 15, 2023AP ! PURDUE PHARMA LP EQ 2MG BASE/2ML (EQ 1MG BASE/ML) A212955 001 Feb 08, 2022

! CHENGDU SHUODE EQ 0.1MG BASE/ML (EQ 0.1MG BASE/ML) A216007 001 Nov 15, 2023

SOLUTION;INTRAMUSCULAR, SUBCUTANEOUS

ZURNAI (AUTOINJECTOR)

+! PURDUE PHARMA LP EQ 1.5MG BASE/0.5ML (EQ 1.5MG BASE/0.5ML) N218590 001 Aug 07, 2024

SPRAY;NASAL

OPVEE

+! INDIVIOR EQ 2.7MG BASE/SPRAY N217470 001 May 22, 2023

NALOXEGOL OXALATE

TABLET;ORAL

MOVANTIK

+ VALINOR EQ 12.5MG BASE N204760 001 Sep 16, 2014

+! EQ 25MG BASE N204760 002 Sep 16, 2014

NALOXONE HYDROCHLORIDE

INJECTABLE;INJECTION

NALOXONEAP HIKMA 0.4MG/ML A070299 001 Sep 24, 1986NALOXONE HYDROCHLORIDEAP ACCORD HLTHCARE 0.4MG/ML A216624 001 Oct 26, 2022AP 1MG/ML A216624 002 Oct 26, 2022AP BAXTER HLTHCARE 0.4MG/ML A214785 001 Jan 29, 2021

CORP

AP 0.4MG/ML A214792 001 Nov 07, 2022AP BPI LABS 1MG/ML A216977 001 Oct 06, 2023AP DR REDDYS 1MG/ML A213209 001 Mar 16, 2020AP ENDO OPERATIONS 1MG/ML A215964 001 Jul 29, 2022AP EUGIA PHARMA 1MG/ML A213279 001 Jan 14, 2021AP 0.4MG/ML A212456 001 Nov 04, 2019AP FRESENIUS KABI USA 0.4MG/ML A213573 001 Apr 02, 2024AP 1MG/ML A213573 002 Apr 02, 2024AP HIKMA 0.4MG/ML A212300 002 Mar 07, 2024AP 1MG/ML A212300 001 Jun 10, 2022AP ! HOSPIRA 0.4MG/ML A070172 001 Sep 24, 1986AP ! 0.4MG/ML A070256 001 Jan 07, 1987AP ! 0.4MG/ML A070257 001 Jan 07, 1987AP ! INTL MEDICATION 1MG/ML A072076 001 Mar 24, 1988AP MANKIND PHARMA 0.4MG/ML A218404 001 Feb 29, 2024AP MYLAN INSTITUTIONAL 0.4MG/ML A204997 001 Mar 06, 2014AP 0.4MG/ML A205014 001 Jun 29, 2016AP MYLAN LABS LTD 1MG/ML A213843 001 Jun 09, 2022AP RISING 0.4MG/ML A208871 001 Feb 28, 2017AP 0.4MG/ML A208872 001 Mar 14, 2017AP SOMERSET THERAPS 0.4MG/ML A207633 001 Aug 08, 2017

LLC

AP 0.4MG/ML A207634 001 Jul 26, 2017

SOLUTION;INTRAMUSCULAR, SUBCUTANEOUS

ZIMHI

+! ZMI PHARMA 5MG/0.5ML (5MG/0.5ML) N212854 001 Oct 15, 2021

SPRAY;NASAL

KLOXXADO

+! HIKMA 8MG/SPRAY N212045 001 Apr 29, 2021

REZENOPY

+! SUMMIT BIOSCI 10MG/SPRAY N215487 001 Apr 19, 2024

SPRAY, METERED;NASAL

REXTOVY

+! AMPHASTAR PHARMS 4MG/SPRAY N208969 001 Mar 07, 2023

INC

PRESCRIPTION DRUG PRODUCT LIST

NALOXONE HYDROCHLORIDE; PENTAZOCINE HYDROCHLORIDE

TABLET; ORAL

NALOXONE HYDROCHLORIDE AND PENTAZOCINE HYDROCHLORIDE

<u>AB</u>	LUPIN	<u>EQ 0.5MG BASE;EQ 50MG BASE</u>	<u>A075735 001</u>	Jul 11, 2001
<u>AB</u>	SUN PHARM INDS LTD	<u>EQ 0.5MG BASE;EQ 50MG BASE</u>	<u>A075523 001</u>	Mar 17, 2000
<u>AB</u>	! WATSON LABS	<u>EQ 0.5MG BASE;EQ 50MG BASE</u>	<u>A074736 001</u>	Jan 21, 1997

NALTREXONE

FOR SUSPENSION, EXTENDED RELEASE; INTRAMUSCULAR

VIVITROL

+!	ALKERMES	380MG/VIAL	N021897 001	Apr 13, 2006
----	----------	------------	-------------	--------------

NALTREXONE HYDROCHLORIDE

TABLET; ORAL

NALTREXONE HYDROCHLORIDE

<u>AB</u>	ACCORD HLTHCARE	<u>50MG</u>	<u>A091205 001</u>	Aug 17, 2011
<u>AB</u>	BARR	<u>50MG</u>	<u>A074918 001</u>	May 08, 1998
<u>AB</u>	CHARTWELL	<u>50MG</u>	<u>A207905 001</u>	Jul 21, 2017
<u>AB</u>	ELITE LABS	<u>50MG</u>	<u>A075274 001</u>	May 26, 1999
<u>AB</u>	! SPECGX LLC	<u>50MG</u>	<u>A076264 002</u>	Mar 22, 2002
<u>AB</u>	SUN PHARM	<u>50MG</u>	<u>A090356 001</u>	Feb 24, 2012
	SPECGX LLC	25MG	A076264 001	Mar 22, 2002
		100MG	A076264 003	Mar 22, 2002

NAPROXEN

SUSPENSION; ORAL

NAPROSYN

<u>AB</u>	+ ATNAHS PHARMA US	<u>25MG/ML</u>	<u>N018965 001</u>	Mar 23, 1987
-----------	--------------------	----------------	--------------------	--------------

NAPROXEN

<u>AB</u>	AMNEAL	<u>25MG/ML</u>	<u>A212705 001</u>	Jul 31, 2020
<u>AB</u>	HETERO LABS LTD III	<u>25MG/ML</u>	<u>A215776 001</u>	Jun 07, 2022
<u>AB</u>	! HIKMA	<u>25MG/ML</u>	<u>A074190 001</u>	Mar 30, 1994
<u>AB</u>	NOVITIUM PHARMA	<u>25MG/ML</u>	<u>A211910 001</u>	Mar 10, 2021

TABLET; ORAL

NAPROSYN

<u>AB</u>	+! ATNAHS PHARMA US	<u>500MG</u>	<u>N017581 004</u>	Apr 15, 1982
-----------	---------------------	--------------	--------------------	--------------

NAPROXEN

<u>AB</u>	ADAPTIS	<u>250MG</u>	<u>A091416 001</u>	Feb 14, 2011
<u>AB</u>		<u>375MG</u>	<u>A091416 002</u>	Feb 14, 2011
<u>AB</u>		<u>500MG</u>	<u>A091416 003</u>	Feb 14, 2011
<u>AB</u>	AMNEAL PHARMS NY	<u>250MG</u>	<u>A075927 001</u>	Dec 18, 2001
<u>AB</u>		<u>375MG</u>	<u>A075927 002</u>	Dec 18, 2001
<u>AB</u>		<u>500MG</u>	<u>A075927 003</u>	Dec 18, 2001
<u>AB</u>	AUROBINDO PHARMA	<u>250MG</u>	<u>A200429 001</u>	Nov 08, 2011
<u>AB</u>		<u>375MG</u>	<u>A200429 002</u>	Nov 08, 2011
<u>AB</u>		<u>500MG</u>	<u>A200429 003</u>	Nov 08, 2011
<u>AB</u>	GLENMARK PHARMS LTD	<u>250MG</u>	<u>A078250 001</u>	Mar 28, 2007
<u>AB</u>		<u>375MG</u>	<u>A078250 002</u>	Mar 28, 2007
<u>AB</u>		<u>500MG</u>	<u>A078250 003</u>	Mar 28, 2007
<u>AB</u>	GRANULES	<u>250MG</u>	<u>A074140 001</u>	Dec 21, 1993
<u>AB</u>		<u>375MG</u>	<u>A074140 002</u>	Dec 21, 1993
<u>AB</u>		<u>500MG</u>	<u>A074140 003</u>	Dec 21, 1993
<u>AB</u>	SCIEGEN PHARMS INC	<u>250MG</u>	<u>A212517 001</u>	Feb 21, 2020
<u>AB</u>		<u>375MG</u>	<u>A212517 002</u>	Feb 21, 2020
<u>AB</u>		<u>500MG</u>	<u>A212517 003</u>	Feb 21, 2020
<u>AB</u>	TEVA	<u>250MG</u>	<u>A074201 001</u>	Dec 21, 1993
<u>AB</u>		<u>375MG</u>	<u>A074201 002</u>	Dec 21, 1993
<u>AB</u>		<u>500MG</u>	<u>A074201 003</u>	Dec 21, 1993
<u>AB</u>	ZYDUS PHARMS USA	<u>250MG</u>	<u>A078620 001</u>	Jun 07, 2007
<u>AB</u>		<u>375MG</u>	<u>A078620 002</u>	Jun 07, 2007
<u>AB</u>		<u>500MG</u>	<u>A078620 003</u>	Jun 07, 2007

TABLET, DELAYED RELEASE; ORAL

EC-NAPROSYN

<u>AB</u>	+! ATNAHS PHARMA US	<u>375MG</u>	<u>N020067 002</u>	Oct 14, 1994
<u>AB</u>	+!	<u>500MG</u>	<u>N020067 003</u>	Oct 14, 1994

NAPROXEN

<u>AB</u>	NOVITIUM PHARMA	<u>375MG</u>	<u>A218497 001</u>	Jun 12, 2024
<u>AB</u>		<u>500MG</u>	<u>A218497 002</u>	Jun 12, 2024
<u>AB</u>	NUVO PHARMS INC	<u>375MG</u>	<u>A091432 001</u>	Sep 19, 2011
<u>AB</u>		<u>500MG</u>	<u>A091432 002</u>	Sep 19, 2011
<u>AB</u>	TEVA	<u>375MG</u>	<u>A075227 001</u>	Jun 30, 1998
<u>AB</u>		<u>500MG</u>	<u>A075227 002</u>	Jun 30, 1998
<u>AB</u>	TRUPHARMA	<u>375MG</u>	<u>A216908 001</u>	May 31, 2023
<u>AB</u>		<u>500MG</u>	<u>A216908 002</u>	May 31, 2023

PRESCRIPTION DRUG PRODUCT LIST

NAPROXEN SODIUM

TABLET; ORAL

ANAPROX DS

<u>AB</u>	<u>+</u> !	ATNAHS PHARMA US	<u>EQ 500MG BASE</u>	<u>N018164 003</u>	Sep 30, 1987
-----------	------------	------------------	----------------------	--------------------	--------------

NAPROXEN SODIUM

<u>AB</u>		AUROBINDO PHARMA LTD	<u>EQ 250MG BASE</u>	<u>A200629 001</u>	Oct 31, 2011
-----------	--	----------------------	----------------------	--------------------	--------------

<u>AB</u>			<u>EQ 500MG BASE</u>	<u>A200629 002</u>	Oct 31, 2011
-----------	--	--	----------------------	--------------------	--------------

<u>AB</u>		DR REDDYS LABS LTD	<u>EQ 250MG BASE</u>	<u>A078486 001</u>	Jul 26, 2007
-----------	--	--------------------	----------------------	--------------------	--------------

<u>AB</u>			<u>EQ 500MG BASE</u>	<u>A078486 002</u>	Jul 26, 2007
-----------	--	--	----------------------	--------------------	--------------

<u>AB</u>		GLENMARK PHARMS LTD	<u>EQ 250MG BASE</u>	<u>A078314 001</u>	Apr 27, 2007
-----------	--	---------------------	----------------------	--------------------	--------------

<u>AB</u>			<u>EQ 500MG BASE</u>	<u>A078314 002</u>	Apr 27, 2007
-----------	--	--	----------------------	--------------------	--------------

<u>AB</u>		SCIEGEN PHARMS INC	<u>EQ 250MG BASE</u>	<u>A212199 001</u>	Oct 30, 2019
-----------	--	--------------------	----------------------	--------------------	--------------

<u>AB</u>			<u>EQ 500MG BASE</u>	<u>A212199 002</u>	Oct 30, 2019
-----------	--	--	----------------------	--------------------	--------------

TABLET, EXTENDED RELEASE; ORAL

NAPRELAN

<u>AB</u>	<u>+</u>	TWI PHARMS	<u>EQ 375MG BASE</u>	<u>N020353 001</u>	Jan 05, 1996
-----------	----------	------------	----------------------	--------------------	--------------

<u>AB</u>	<u>+</u>		<u>EQ 500MG BASE</u>	<u>N020353 002</u>	Jan 05, 1996
-----------	----------	--	----------------------	--------------------	--------------

<u>AB</u>	<u>+</u> !		<u>EQ 750MG BASE</u>	<u>N020353 003</u>	Jan 05, 1996
-----------	------------	--	----------------------	--------------------	--------------

NAPROXEN SODIUM

<u>AB</u>		ACTAVIS LABS FL INC	<u>EQ 375MG BASE</u>	<u>A075416 002</u>	Apr 23, 2003
-----------	--	---------------------	----------------------	--------------------	--------------

<u>AB</u>			<u>EQ 500MG BASE</u>	<u>A075416 001</u>	Aug 27, 2002
-----------	--	--	----------------------	--------------------	--------------

<u>AB</u>			<u>EQ 750MG BASE</u>	<u>A075416 003</u>	Aug 11, 2016
-----------	--	--	----------------------	--------------------	--------------

NAPROXEN SODIUM; SUMATRIPTAN SUCCINATE

TABLET; ORAL

SUMATRIPTAN AND NAPROXEN SODIUM

<u>AB</u>		AUROBINDO PHARMA LTD	<u>500MG;EQ 85MG BASE</u>	<u>A207457 001</u>	Feb 15, 2018
-----------	--	----------------------	---------------------------	--------------------	--------------

<u>AB</u>		RISING	<u>500MG;EQ 85MG BASE</u>	<u>A090872 001</u>	Sep 04, 2018
-----------	--	--------	---------------------------	--------------------	--------------

<u>AB</u>		SUN PHARM	<u>500MG;EQ 85MG BASE</u>	<u>A202803 001</u>	Jul 20, 2018
-----------	--	-----------	---------------------------	--------------------	--------------

TREXIMET

<u>AB</u>	<u>+</u> !	CURRAX	<u>500MG;EQ 85MG BASE</u>	<u>N021926 001</u>	Apr 15, 2008
-----------	------------	--------	---------------------------	--------------------	--------------

NARATRIPTAN HYDROCHLORIDE

TABLET; ORAL

NARATRIPTAN

<u>AB</u>		HERITAGE	<u>EQ 1MG BASE</u>	<u>A200502 001</u>	Feb 28, 2011
-----------	--	----------	--------------------	--------------------	--------------

<u>AB</u>	<u>!</u>		<u>EQ 2.5MG BASE</u>	<u>A200502 002</u>	Feb 28, 2011
-----------	----------	--	----------------------	--------------------	--------------

<u>AB</u>		HIKMA	<u>EQ 1MG BASE</u>	<u>A090381 001</u>	Jul 07, 2010
-----------	--	-------	--------------------	--------------------	--------------

<u>AB</u>			<u>EQ 2.5MG BASE</u>	<u>A090381 002</u>	Jul 07, 2010
-----------	--	--	----------------------	--------------------	--------------

<u>AB</u>		ORBION PHARMS	<u>EQ 1MG BASE</u>	<u>A091441 001</u>	Apr 30, 2012
-----------	--	---------------	--------------------	--------------------	--------------

<u>AB</u>			<u>EQ 2.5MG BASE</u>	<u>A091441 002</u>	Apr 30, 2012
-----------	--	--	----------------------	--------------------	--------------

<u>AB</u>		PADAGIS US	<u>EQ 1MG BASE</u>	<u>A091326 001</u>	Jul 08, 2010
-----------	--	------------	--------------------	--------------------	--------------

<u>AB</u>			<u>EQ 2.5MG BASE</u>	<u>A091326 002</u>	Jul 08, 2010
-----------	--	--	----------------------	--------------------	--------------

NATAMYCIN

SUSPENSION; OPHTHALMIC

NATACYN

<u>+</u> !	HARROW EYE	5%	<u>N050514 001</u>
------------	------------	----	--------------------

NATEGLINIDE

TABLET; ORAL

NATEGLINIDE

<u>AB</u>		CADILA PHARMS LTD	<u>60MG</u>	<u>A206432 001</u>	Apr 19, 2019
-----------	--	-------------------	-------------	--------------------	--------------

<u>AB</u>			<u>120MG</u>	<u>A206432 002</u>	Apr 19, 2019
-----------	--	--	--------------	--------------------	--------------

<u>AB</u>		DR REDDYS LABS LTD	<u>60MG</u>	<u>A077461 001</u>	Sep 09, 2009
-----------	--	--------------------	-------------	--------------------	--------------

<u>AB</u>			<u>120MG</u>	<u>A077461 002</u>	Sep 09, 2009
-----------	--	--	--------------	--------------------	--------------

<u>AB</u>		RISING	<u>60MG</u>	<u>A205544 001</u>	Jun 18, 2018
-----------	--	--------	-------------	--------------------	--------------

<u>AB</u>			<u>120MG</u>	<u>A205544 002</u>	Jun 18, 2018
-----------	--	--	--------------	--------------------	--------------

<u>AB</u>		STRIDES PHARMA	<u>60MG</u>	<u>A077463 001</u>	Sep 09, 2009
-----------	--	----------------	-------------	--------------------	--------------

<u>AB</u>			<u>120MG</u>	<u>A077463 002</u>	Sep 09, 2009
-----------	--	--	--------------	--------------------	--------------

<u>AB</u>		WATSON LABS	<u>60MG</u>	<u>A077462 001</u>	Mar 30, 2011
-----------	--	-------------	-------------	--------------------	--------------

<u>AB</u>			<u>120MG</u>	<u>A077462 002</u>	Mar 30, 2011
-----------	--	--	--------------	--------------------	--------------

<u>AB</u>		ZYDUS PHARMS	<u>60MG</u>	<u>A205248 001</u>	Jul 06, 2016
-----------	--	--------------	-------------	--------------------	--------------

<u>AB</u>	<u>!</u>		<u>120MG</u>	<u>A205248 002</u>	Jul 06, 2016
-----------	----------	--	--------------	--------------------	--------------

NEBIVOLOL HYDROCHLORIDE

TABLET; ORAL

BYSTOLIC

<u>AB</u>	<u>+</u>	ALLERGAN	<u>EQ 2.5MG BASE</u>	<u>N021742 002</u>	Dec 17, 2007
-----------	----------	----------	----------------------	--------------------	--------------

<u>AB</u>	<u>+</u>		<u>EQ 5MG BASE</u>	<u>N021742 003</u>	Dec 17, 2007
-----------	----------	--	--------------------	--------------------	--------------

<u>AB</u>	<u>+</u>		<u>EQ 10MG BASE</u>	<u>N021742 004</u>	Dec 17, 2007
-----------	----------	--	---------------------	--------------------	--------------

<u>AB</u>	<u>+</u> !		<u>EQ 20MG BASE</u>	<u>N021742 005</u>	Oct 08, 2008
-----------	------------	--	---------------------	--------------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

NEBIVOLOL HYDROCHLORIDE

TABLET; ORAL

NEBIVOLOL HYDROCHLORIDE

<u>AB</u>	ANI PHARMS	<u>EQ 2.5MG BASE</u>	<u>A203659 001</u>	Apr 16, 2015
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A203659 002</u>	Apr 16, 2015
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A203659 003</u>	Apr 16, 2015
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A203659 004</u>	Apr 16, 2015
<u>AB</u>	AUROBINDO PHARMA LTD	<u>EQ 2.5MG BASE</u>	<u>A211053 001</u>	Dec 17, 2021
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A211053 002</u>	Dec 17, 2021
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A211053 003</u>	Dec 17, 2021
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A211053 004</u>	Dec 17, 2021
<u>AB</u>	BEXIMCO PHARMS USA	<u>EQ 2.5MG BASE</u>	<u>A216568 001</u>	Mar 30, 2023
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A216568 002</u>	Mar 30, 2023
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A216568 003</u>	Mar 30, 2023
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A216568 004</u>	Mar 30, 2023
<u>AB</u>	CADILA PHARMS LTD	<u>EQ 2.5MG BASE</u>	<u>A208717 001</u>	Dec 17, 2021
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A208717 002</u>	Dec 17, 2021
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A208717 003</u>	Dec 17, 2021
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A208717 004</u>	Dec 17, 2021
<u>AB</u>	GLENMARK PHARMS LTD	<u>EQ 2.5MG BASE</u>	<u>A203821 001</u>	May 25, 2017
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A203821 002</u>	May 25, 2017
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A203821 003</u>	May 25, 2017
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A203821 004</u>	May 25, 2017
<u>AB</u>	HETERO LABS LTD III	<u>EQ 2.5MG BASE</u>	<u>A203825 001</u>	Nov 03, 2020
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A203825 002</u>	Nov 03, 2020
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A203825 003</u>	Nov 03, 2020
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A203825 004</u>	Nov 03, 2020
<u>AB</u>	INDCHEMIE HEALTH	<u>EQ 2.5MG BASE</u>	<u>A203828 001</u>	Jul 29, 2015
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A203828 002</u>	Jul 29, 2015
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A203828 003</u>	Jul 29, 2015
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A203828 004</u>	Jul 29, 2015
<u>AB</u>	MANKIND PHARMA	<u>EQ 2.5MG BASE</u>	<u>A216172 001</u>	Nov 14, 2022
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A216172 002</u>	Nov 14, 2022
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A216172 003</u>	Nov 14, 2022
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A216172 004</u>	Nov 14, 2022
<u>AB</u>	MSN	<u>EQ 2.5MG BASE</u>	<u>A217397 001</u>	Oct 01, 2024
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A217397 002</u>	Oct 01, 2024
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A217397 003</u>	Oct 01, 2024
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A217397 004</u>	Oct 01, 2024
<u>AB</u>	PRINSTON INC	<u>EQ 2.5MG BASE</u>	<u>A212682 001</u>	Apr 07, 2022
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A212682 002</u>	Apr 07, 2022
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A212682 003</u>	Apr 07, 2022
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A212682 004</u>	Apr 07, 2022
<u>AB</u>	REYOUNG	<u>EQ 2.5MG BASE</u>	<u>A212917 001</u>	Dec 17, 2021
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A212917 002</u>	Dec 17, 2021
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A212917 003</u>	Dec 17, 2021
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A212917 004</u>	Dec 17, 2021
<u>AB</u>	TORRENT	<u>EQ 2.5MG BASE</u>	<u>A203966 001</u>	Mar 02, 2018
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A203966 002</u>	Mar 02, 2018
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A203966 003</u>	Mar 02, 2018
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A203966 004</u>	Mar 02, 2018
<u>AB</u>	UNICHEM	<u>EQ 2.5MG BASE</u>	<u>A213830 001</u>	Mar 15, 2022
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A213830 002</u>	Mar 15, 2022
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A213830 003</u>	Mar 15, 2022
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A213830 004</u>	Mar 15, 2022

NEBIVOLOL HYDROCHLORIDE; VALSARTAN

TABLET; ORAL

VYDUO

PRINSTON INC EQ 5MG BASE; 80MG A210596 001 Sep 19, 2022

NEDOSIRAN SODIUM

SOLUTION; INJECTION

RIVFLOZA

+	NOVO	EQ 80MG BASE/0.5ML (EQ 160MG BASE/ML)	N215842 001	Sep 29, 2023
+		EQ 128MG BASE/0.8ML (EQ 160MG BASE/ML)	N215842 002	Sep 29, 2023
+		EQ 160MG BASE/ML (EQ 160MG BASE/ML)	N215842 003	Sep 29, 2023

PRESCRIPTION DRUG PRODUCT LIST

NEFAZODONE HYDROCHLORIDE

TABLET; ORAL

NEFAZODONE HYDROCHLORIDE

TEVA	50MG	A076037	001	Sep 16, 2003
	100MG	A076037	002	Sep 16, 2003
	150MG	A076037	003	Sep 16, 2003
	200MG	A076037	004	Sep 16, 2003
!	250MG	A076037	005	Sep 16, 2003

NELARABINE

INJECTABLE; INTRAVENOUS

ARRANON

AP	+	SANDOZ	<u>250MG/50ML (5MG/ML)</u>	<u>N021877</u>	<u>001</u>	Oct 28, 2005
----	---	--------	----------------------------	----------------	------------	--------------

NELARABINE

AP		ALEMBIC	<u>250MG/50ML (5MG/ML)</u>	<u>A218554</u>	<u>001</u>	Aug 01, 2024
AP		AMNEAL	<u>250MG/50ML (5MG/ML)</u>	<u>A216346</u>	<u>001</u>	Apr 04, 2023
AP		DR REDDYS	<u>250MG/50ML (5MG/ML)</u>	<u>A216934</u>	<u>001</u>	Dec 23, 2022
AP		GLAND PHARMA LTD	<u>250MG/50ML (5MG/ML)</u>	<u>A212605</u>	<u>001</u>	Jan 03, 2024
AP		MEITHEAL	<u>250MG/50ML (5MG/ML)</u>	<u>A216038</u>	<u>001</u>	Jan 10, 2023
AP		MSN	<u>250MG/50ML (5MG/ML)</u>	<u>A216948</u>	<u>001</u>	Sep 13, 2024
AP		NEXUS	<u>250MG/50ML (5MG/ML)</u>	<u>A215057</u>	<u>001</u>	Jun 02, 2023
AP		SHORLA	<u>250MG/50ML (5MG/ML)</u>	<u>A214809</u>	<u>001</u>	Mar 03, 2023
AP		XGEN PHARMS	<u>250MG/50ML (5MG/ML)</u>	<u>A216510</u>	<u>001</u>	Sep 05, 2024
AP		ZYDUS PHARMS	<u>250MG/50ML (5MG/ML)</u>	<u>A215037</u>	<u>001</u>	Nov 17, 2021

NELFINAVIR MESYLATE

TABLET; ORAL

VIRACEPT

+	!	AGOURON PHARMS	EQ 250MG BASE	N020779	001	Mar 14, 1997
+	!		EQ 625MG BASE	N021503	001	Apr 30, 2003

NEOMYCIN SULFATE

TABLET; ORAL

NEOMYCIN SULFATE

AA		ADRASTEIA PHARMA	<u>500MG</u>	<u>A065220</u>	<u>001</u>	Jul 28, 2006
AA	!	TEVA	<u>500MG</u>	<u>A060304</u>	<u>001</u>	

NEOMYCIN SULFATE; POLYMYXIN B SULFATE

SOLUTION; IRRIGATION

NEOMYCIN AND POLYMYXIN B SULFATE

AT		WATSON LABS	<u>EQ 40MG BASE/ML; 200,000 UNITS/ML</u>	<u>A062664</u>	<u>001</u>	Apr 08, 1986
AT		XGEN PHARMS	<u>EQ 40MG BASE/ML; 200,000 UNITS/ML</u>	<u>A065106</u>	<u>001</u>	Jan 31, 2006
AT			<u>EQ 40MG BASE/ML; 200,000 UNITS/ML</u>	<u>A065108</u>	<u>001</u>	Jan 31, 2006

NEOSPORIN G.U. IRRIGANT

AT	!	MONARCH PHARMS	<u>EQ 40MG BASE/ML; 200,000 UNITS/ML</u>	<u>A060707</u>	<u>001</u>	
----	---	----------------	--	----------------	------------	--

NEOSTIGMINE METHYLSULFATE

SOLUTION; INTRAVENOUS

BLOXIVERZ

AP	+	EXELA PHARMA	<u>5MG/10ML (0.5MG/ML)</u>	<u>N204078</u>	<u>001</u>	May 31, 2013
AP	+		<u>10MG/10ML (1MG/ML)</u>	<u>N204078</u>	<u>002</u>	May 31, 2013

NEOSTIGMINE METHYLSULFATE

AP		ADAPTIS	<u>5MG/10ML (0.5MG/ML)</u>	<u>A210652</u>	<u>001</u>	Oct 01, 2021
AP			<u>10MG/10ML (1MG/ML)</u>	<u>A210652</u>	<u>002</u>	Oct 01, 2021
AP		AMNEAL	<u>5MG/10ML (0.5MG/ML)</u>	<u>A210051</u>	<u>001</u>	Jun 15, 2018
AP			<u>10MG/10ML (1MG/ML)</u>	<u>A210051</u>	<u>002</u>	Jun 15, 2018
AP		AMPHASTAR PHARMS INC	<u>5MG/10ML (0.5MG/ML)</u>	<u>A209933</u>	<u>001</u>	Sep 25, 2017
AP			<u>10MG/10ML (1MG/ML)</u>	<u>A209933</u>	<u>002</u>	Sep 25, 2017
AP		AMRING PHARMS	<u>5MG/10ML (0.5MG/ML)</u>	<u>A210989</u>	<u>001</u>	Aug 22, 2018
AP			<u>10MG/10ML (1MG/ML)</u>	<u>A210989</u>	<u>002</u>	Aug 22, 2018
AP		BE PHARMS	<u>5MG/10ML (0.5MG/ML)</u>	<u>A212512</u>	<u>001</u>	May 13, 2019
AP			<u>10MG/10ML (1MG/ML)</u>	<u>A212512</u>	<u>002</u>	May 13, 2019
AP		CAPLIN	<u>5MG/10ML (0.5MG/ML)</u>	<u>A213074</u>	<u>001</u>	Apr 20, 2021
AP			<u>10MG/10ML (1MG/ML)</u>	<u>A213074</u>	<u>002</u>	Apr 20, 2021
AP		DR REDDYS	<u>3MG/3ML (1MG/ML)</u>	<u>A216291</u>	<u>001</u>	Jul 06, 2022
AP			<u>5MG/10ML (0.5MG/ML)</u>	<u>A209135</u>	<u>001</u>	Jul 10, 2018
AP			<u>10MG/10ML (1MG/ML)</u>	<u>A209135</u>	<u>002</u>	Jul 10, 2018
AP		ENDO OPERATIONS	<u>5MG/10ML (0.5MG/ML)</u>	<u>A208405</u>	<u>001</u>	Apr 26, 2017
AP			<u>10MG/10ML (1MG/ML)</u>	<u>A208405</u>	<u>002</u>	Apr 26, 2017
AP	+	FRESENIUS KABI USA	<u>3MG/3ML (1MG/ML)</u>	<u>N203629</u>	<u>003</u>	Sep 18, 2018
AP		GLAND PHARMA LTD	<u>5MG/10ML (0.5MG/ML)</u>	<u>A212968</u>	<u>001</u>	Oct 16, 2019
AP			<u>10MG/10ML (1MG/ML)</u>	<u>A212968</u>	<u>002</u>	Oct 16, 2019
AP		HIKMA	<u>3MG/3ML (1MG/ML)</u>	<u>A216206</u>	<u>001</u>	Jul 13, 2022
AP			<u>5MG/10ML (0.5MG/ML)</u>	<u>A207042</u>	<u>001</u>	Dec 28, 2015

PRESCRIPTION DRUG PRODUCT LIST

NEOSTIGMINE METHYLSULFATE

SOLUTION; INTRAVENOUS

NEOSTIGMINE METHYLSULFATE

<u>AP</u>		<u>10MG/10ML (1MG/ML)</u>	<u>A207042 002</u>	Dec 28, 2015
<u>AP</u>	MEITHEAL	<u>5MG/10ML (0.5MG/ML)</u>	<u>A212804 001</u>	Apr 05, 2021
<u>AP</u>		<u>10MG/10ML (1MG/ML)</u>	<u>A212804 002</u>	Apr 05, 2021
<u>AP</u>	SAGENT PHARMS INC	<u>5MG/10ML (0.5MG/ML)</u>	<u>A216542 001</u>	Feb 17, 2023
<u>AP</u>		<u>10MG/10ML (1MG/ML)</u>	<u>A216542 002</u>	Feb 17, 2023
<u>AP</u>	UMEDICA	<u>5MG/10ML (0.5MG/ML)</u>	<u>A212627 001</u>	Nov 03, 2022
<u>AP</u>		<u>10MG/10ML (1MG/ML)</u>	<u>A212627 002</u>	Nov 03, 2022

NEOSTIGMINE METHYLSULFATE

<u>AP</u>	AVET LIFESCIENCES	<u>5MG/10ML (0.5MG/ML)</u>	<u>A208230 001</u>	Nov 25, 2022
<u>AP</u>		<u>10MG/10ML (1MG/ML)</u>	<u>A208230 002</u>	Nov 25, 2022
	BLOXIVERZ			
	+! EXELA PHARMA	5MG/5ML (1MG/ML)	N204078 003	Oct 27, 2023
	NEOSTIGMINE METHYLSULFATE			
	+! FRESENIUS KABI USA	5MG/10ML (0.5MG/ML)	N203629 001	Jan 08, 2015
	+!	10MG/10ML (1MG/ML)	N203629 002	Jan 08, 2015

NEPAFENAC

SUSPENSION/DROPS; OPHTHALMIC

ILEVRO

+! HARROW EYE

0.3%

N203491 001 Oct 16, 2012

NEVANAC

+! HARROW EYE

0.1%

N021862 001 Aug 19, 2005

NERATINIB MALEATE

TABLET; ORAL

NERLYNX

+! PUMA BIOTECH

EQ 40MG BASE

N208051 001 Jul 17, 2017

NETARSUDIL MESYLATE

SOLUTION/DROPS; OPHTHALMIC

RHOPRESSA

+! ALCON LABS INC

EQ 0.02% BASE

N208254 001 Dec 18, 2017

NETUPITANT; PALONOSETRON HYDROCHLORIDE

CAPSULE; ORAL

AKYNZEO

+! HELSINN HLTHCARE

300MG; EQ 0.5MG BASE

N205718 001 Oct 10, 2014

NEVIRAPINE

SUSPENSION; ORAL

NEVIRAPINE

<u>AA</u>	! AUROBINDO	<u>50MG/5ML</u>	<u>A077702 001</u>	May 22, 2012
<u>AA</u>	CIPLA	<u>50MG/5ML</u>	<u>A207684 001</u>	Aug 03, 2017

TABLET; ORAL

NEVIRAPINE

<u>AB</u>	AUROBINDO	<u>200MG</u>	<u>A077521 001</u>	May 22, 2012
<u>AB</u>	CIPLA	<u>200MG</u>	<u>A077956 001</u>	May 22, 2012
<u>AB</u>	HETERO LABS LTD III	<u>200MG</u>	<u>A078584 001</u>	May 22, 2012
<u>AB</u>	! MACLEODS PHARMS LTD	<u>200MG</u>	<u>A090688 001</u>	Jan 14, 2019
<u>AB</u>	MICRO LABS LTD	<u>200MG</u>	<u>A203080 001</u>	May 22, 2012
<u>AB</u>	MYLAN PHARMS INC	<u>200MG</u>	<u>A202523 001</u>	May 22, 2012
<u>AB</u>	STRIDES PHARMA	<u>200MG</u>	<u>A078195 001</u>	May 22, 2012

TABLET, EXTENDED RELEASE; ORAL

NEVIRAPINE

<u>AB</u>	! MACLEODS PHARMS LTD	<u>400MG</u>	<u>A206879 001</u>	Oct 06, 2017
<u>AB</u>	MYLAN	<u>400MG</u>	<u>A205651 001</u>	Oct 27, 2014
<u>AB</u>	SANDOZ	<u>400MG</u>	<u>A203411 001</u>	Apr 03, 2014

NIACIN

TABLET; ORAL

NIACOR

! AVONDALE PHARMS

500MG

A040378 001 May 03, 2000

TABLET, EXTENDED RELEASE; ORAL

NIACIN

<u>AB</u>	AMNEAL PHARMS	<u>500MG</u>	<u>A203578 001</u>	Jul 24, 2015
<u>AB</u>		<u>750MG</u>	<u>A204178 001</u>	Dec 11, 2015
<u>AB</u>		<u>1GM</u>	<u>A203578 002</u>	Jul 24, 2015
<u>AB</u>	AUROBINDO PHARMA LTD	<u>500MG</u>	<u>A209236 001</u>	Feb 01, 2018
<u>AB</u>		<u>750MG</u>	<u>A209236 002</u>	Feb 01, 2018
<u>AB</u>		<u>1GM</u>	<u>A209236 003</u>	Feb 01, 2018
<u>AB</u>	BARR	<u>500MG</u>	<u>A076378 001</u>	Apr 26, 2005
<u>AB</u>		<u>750MG</u>	<u>A076378 002</u>	Apr 26, 2005

PRESCRIPTION DRUG PRODUCT LIST

NIACIN

TABLET, EXTENDED RELEASE;ORAL

NIACIN

<u>AB</u>		<u>1GM</u>	<u>A076250 001</u>	Apr 14, 2005
<u>AB</u>	CHARTWELL RX	<u>500MG</u>	<u>A090860 001</u>	Mar 20, 2014
<u>AB</u>		<u>750MG</u>	<u>A090892 001</u>	Mar 20, 2014
<u>AB</u>		<u>1GM</u>	<u>A090446 001</u>	Mar 20, 2014
<u>AB</u>	HIBROW HLTHCARE	<u>500MG</u>	<u>A213090 001</u>	Aug 25, 2023
<u>AB</u>		<u>750MG</u>	<u>A213090 002</u>	Aug 25, 2023
<u>AB</u>		<u>1GM</u>	<u>A213090 003</u>	Aug 25, 2023
<u>AB</u>	LANNETT CO INC	<u>500MG</u>	<u>A203899 001</u>	Jun 16, 2017
<u>AB</u>		<u>1GM</u>	<u>A203899 002</u>	Jun 16, 2017
<u>AB</u>	MACLEODS PHARMS LTD	<u>500MG</u>	<u>A204934 001</u>	Mar 03, 2022
<u>AB</u>		<u>1GM</u>	<u>A204934 002</u>	Mar 03, 2022
<u>AB</u>	SUN PHARM	<u>500MG</u>	<u>A200484 001</u>	Apr 23, 2014
<u>AB</u>	!	<u>750MG</u>	<u>A201273 001</u>	Apr 23, 2014
<u>AB</u>	!	<u>1GM</u>	<u>A200484 002</u>	Apr 23, 2014

NICARDIPINE HYDROCHLORIDE

CAPSULE;ORAL

NICARDIPINE HYDROCHLORIDE

<u>AB</u>	ANI PHARMS	<u>20MG</u>	<u>A074670 001</u>	Oct 28, 1996
<u>AB</u>		<u>30MG</u>	<u>A074670 002</u>	Oct 28, 1996
<u>AB</u>	BIONPHARMA	<u>20MG</u>	<u>A217555 001</u>	May 03, 2023
<u>AB</u>		<u>30MG</u>	<u>A217555 002</u>	May 03, 2023
<u>AB</u>	EPIC PHARMA LLC	<u>20MG</u>	<u>A074928 001</u>	Mar 19, 1998
<u>AB</u>	!	<u>30MG</u>	<u>A074928 002</u>	Mar 19, 1998
<u>AB</u>	GLENMARK PHARMS LTD	<u>20MG</u>	<u>A216357 001</u>	Dec 16, 2022
<u>AB</u>		<u>30MG</u>	<u>A216357 002</u>	Dec 16, 2022
<u>AB</u>	SENORES PHARMS	<u>20MG</u>	<u>A215377 001</u>	Jul 17, 2023
<u>AB</u>		<u>30MG</u>	<u>A215377 002</u>	Jul 17, 2023
<u>AB</u>	SKG PHARMA	<u>20MG</u>	<u>A218202 001</u>	Apr 10, 2024
<u>AB</u>		<u>30MG</u>	<u>A218202 002</u>	Apr 10, 2024

INJECTABLE;INJECTION

NICARDIPINE HYDROCHLORIDE

<u>AP</u>	!	AM REGENT	<u>25MG/10ML (2.5MG/ML)</u>	<u>A090534 001</u>	Nov 17, 2009
<u>AP</u>		AMNEAL	<u>25MG/10ML (2.5MG/ML)</u>	<u>A215406 001</u>	Oct 31, 2024
<u>AP</u>		CHENGDU SHUODE	<u>25MG/10ML (2.5MG/ML)</u>	<u>A217548 001</u>	Sep 06, 2024
<u>AP</u>		EUGIA PHARMA	<u>25MG/10ML (2.5MG/ML)</u>	<u>A211121 001</u>	Apr 08, 2021
<u>AP</u>	+	HIKMA INTL PHARMS	<u>25MG/10ML (2.5MG/ML)</u>	<u>N022276 001</u>	Jul 24, 2008
<u>AP</u>		MICRO LABS	<u>25MG/10ML (2.5MG/ML)</u>	<u>A216420 001</u>	Jul 07, 2023
<u>AP</u>		RK PHARMA	<u>25MG/10ML (2.5MG/ML)</u>	<u>A090664 001</u>	Nov 17, 2009
<u>AP</u>		WOCKHARDT	<u>25MG/10ML (2.5MG/ML)</u>	<u>A090671 001</u>	Nov 17, 2009

INJECTABLE;INTRAVENOUS

CARDENE IN 0.83% SODIUM CHLORIDE IN PLASTIC CONTAINER

<u>AP</u>	+	CHIESI	<u>40MG/200ML (0.2MG/ML)</u>	<u>N019734 004</u>	Nov 07, 2008
-----------	---	--------	------------------------------	--------------------	--------------

CARDENE IN 0.86% SODIUM CHLORIDE IN PLASTIC CONTAINER

<u>AP</u>	+	CHIESI	<u>20MG/200ML (0.1MG/ML)</u>	<u>N019734 003</u>	Jul 31, 2008
-----------	---	--------	------------------------------	--------------------	--------------

NICARDIPINE HYDROCHLORIDE IN 0.83% SODIUM CHLORIDE

<u>AP</u>		INFORLIFE	<u>40MG/200ML (0.2MG/ML)</u>	<u>A203978 002</u>	Apr 17, 2024
-----------	--	-----------	------------------------------	--------------------	--------------

NICARDIPINE HYDROCHLORIDE IN 0.86% SODIUM CHLORIDE

<u>AP</u>		INFORLIFE	<u>20MG/200ML (0.1MG/ML)</u>	<u>A203978 001</u>	Apr 17, 2024
-----------	--	-----------	------------------------------	--------------------	--------------

NICARDIPINE HYDROCHLORIDE IN 0.9% SODIUM CHLORIDE

<u>AP</u>		CIPLA	<u>20MG/200ML (0.1MG/ML)</u>	<u>A215592 001</u>	Sep 24, 2024
-----------	--	-------	------------------------------	--------------------	--------------

<u>AP</u>			<u>40MG/200ML (0.2MG/ML)</u>	<u>A215592 002</u>	Sep 24, 2024
-----------	--	--	------------------------------	--------------------	--------------

<u>AP</u>	+	HIKMA INTL PHARMS	<u>20MG/200ML (0.1MG/ML)</u>	<u>N022276 002</u>	Apr 07, 2016
-----------	---	-------------------	------------------------------	--------------------	--------------

<u>AP</u>	+		<u>40MG/200ML (0.2MG/ML)</u>	<u>N022276 003</u>	Apr 07, 2016
-----------	---	--	------------------------------	--------------------	--------------

CARDENE IN 4.8% DEXTROSE IN PLASTIC CONTAINER

	+	CHIESI	20MG/200ML (0.1MG/ML)	N019734 002	Jul 31, 2008
--	---	--------	-----------------------	-------------	--------------

NICOTINE

SPRAY, METERED;NASAL

NICOTROL

	+	PFIZER INC	0.5MG/SPRAY	N020385 001	Mar 22, 1996
--	---	------------	-------------	-------------	--------------

NIFEDIPINE

CAPSULE;ORAL

NIFEDIPINE

<u>AB</u>	ACELLA	<u>10MG</u>	<u>A072781 001</u>	Jul 30, 1993
<u>AB</u>	ACTAVIS ELIZABETH	<u>10MG</u>	<u>A072579 001</u>	Jan 08, 1991
<u>AB</u>	HERITAGE PHARMA	<u>10MG</u>	<u>A202644 001</u>	Apr 25, 2013
<u>AB</u>		<u>20MG</u>	<u>A202644 002</u>	Apr 25, 2013
<u>AB</u>	VELZEN PHARMA PVT	<u>10MG</u>	<u>A073250 001</u>	Oct 08, 1991
<u>AB</u>		<u>20MG</u>	<u>A074045 001</u>	Apr 30, 1992

PRESCRIPTION DRUG PRODUCT LIST

NIFEDIPINE

CAPSULE; ORAL

PROCARDIA

<u>AB</u>	<u>+</u> !	PFIZER	<u>10MG</u>	<u>N018482</u>	<u>001</u>		
-----------	------------	--------	-------------	----------------	------------	--	--

TABLET, EXTENDED RELEASE; ORAL

NIFEDIPINE

<u>AB1</u>		AUROBINDO PHARMA	<u>30MG</u>	<u>A213361</u>	<u>001</u>	Jul 19, 2021	
<u>AB1</u>			<u>60MG</u>	<u>A213361</u>	<u>002</u>	Jul 19, 2021	
<u>AB1</u>			<u>90MG</u>	<u>A213361</u>	<u>003</u>	Jul 19, 2021	
<u>AB1</u>		NOVAST LABS	<u>30MG</u>	<u>A202987</u>	<u>001</u>	Aug 25, 2016	
<u>AB1</u>	<u>!</u>		<u>60MG</u>	<u>A202987</u>	<u>002</u>	Aug 25, 2016	
<u>AB1</u>	<u>!</u>		<u>90MG</u>	<u>A202987</u>	<u>003</u>	Aug 25, 2016	
<u>AB1</u>		VALEANT PHARMS NORTH	<u>30MG</u>	<u>A075269</u>	<u>001</u>	Dec 04, 2000	
<u>AB1</u>			<u>60MG</u>	<u>A075269</u>	<u>002</u>	Dec 04, 2000	
<u>AB1</u>			<u>90MG</u>	<u>A076070</u>	<u>001</u>	Aug 16, 2002	
<u>AB1</u>		ZYDUS PHARMS	<u>30MG</u>	<u>A210184</u>	<u>001</u>	Jun 29, 2018	
<u>AB1</u>			<u>60MG</u>	<u>A210184</u>	<u>002</u>	Jun 29, 2018	
<u>AB1</u>			<u>90MG</u>	<u>A210184</u>	<u>003</u>	Jun 29, 2018	
<u>AB2</u>		ALEMBIC	<u>30MG</u>	<u>A216896</u>	<u>001</u>	Nov 18, 2022	
<u>AB2</u>			<u>60MG</u>	<u>A216896</u>	<u>002</u>	Nov 18, 2022	
<u>AB2</u>			<u>90MG</u>	<u>A216896</u>	<u>003</u>	Nov 18, 2022	
<u>AB2</u>		ALKEM LABS LTD	<u>30MG</u>	<u>A216067</u>	<u>001</u>	Mar 29, 2022	
<u>AB2</u>			<u>60MG</u>	<u>A216067</u>	<u>002</u>	Mar 29, 2022	
<u>AB2</u>			<u>90MG</u>	<u>A216067</u>	<u>003</u>	Mar 29, 2022	
<u>AB2</u>		ELITE PHARM SOLUTION	<u>90MG</u>	<u>A212016</u>	<u>001</u>	Nov 18, 2020	
<u>AB2</u>		NOVAST LABS	<u>30MG</u>	<u>A210614</u>	<u>001</u>	Mar 12, 2019	
<u>AB2</u>			<u>60MG</u>	<u>A210614</u>	<u>002</u>	Mar 12, 2019	
<u>AB2</u>			<u>90MG</u>	<u>A210614</u>	<u>003</u>	Mar 12, 2019	
<u>AB2</u>		OSMOTICA PHARM US	<u>30MG</u>	<u>A077127</u>	<u>001</u>	Nov 21, 2005	
<u>AB2</u>			<u>60MG</u>	<u>A077127</u>	<u>002</u>	Nov 21, 2005	
<u>AB2</u>			<u>90MG</u>	<u>A077127</u>	<u>003</u>	Oct 03, 2007	
<u>AB2</u>		SPIL	<u>30MG</u>	<u>A210838</u>	<u>001</u>	Apr 16, 2019	
<u>AB2</u>			<u>60MG</u>	<u>A210838</u>	<u>002</u>	Apr 16, 2019	
<u>AB2</u>			<u>90MG</u>	<u>A210838</u>	<u>003</u>	Apr 16, 2019	
<u>AB2</u>		TWI PHARMS	<u>30MG</u>	<u>A203126</u>	<u>001</u>	Apr 03, 2014	
<u>AB2</u>			<u>60MG</u>	<u>A203126</u>	<u>002</u>	Apr 03, 2014	
<u>AB2</u>			<u>90MG</u>	<u>A203126</u>	<u>003</u>	Apr 03, 2014	
<u>AB2</u>		VALEANT PHARMS NORTH	<u>30MG</u>	<u>A075289</u>	<u>002</u>	Feb 06, 2001	
<u>AB2</u>			<u>60MG</u>	<u>A075289</u>	<u>001</u>	Sep 27, 2000	
<u>AB2</u>		ZYDUS PHARMS	<u>30MG</u>	<u>A210012</u>	<u>001</u>	Dec 19, 2017	
<u>AB2</u>			<u>60MG</u>	<u>A210012</u>	<u>002</u>	Dec 19, 2017	
<u>AB2</u>			<u>90MG</u>	<u>A210012</u>	<u>003</u>	Dec 19, 2017	
		<u>PROCARDIA XL</u>					
<u>AB2</u>	<u>+</u>	PFIZER	<u>30MG</u>	<u>N019684</u>	<u>001</u>	Sep 06, 1989	
<u>AB2</u>	<u>+</u>		<u>60MG</u>	<u>N019684</u>	<u>002</u>	Sep 06, 1989	
<u>AB2</u>	<u>+</u> !		<u>90MG</u>	<u>N019684</u>	<u>003</u>	Sep 06, 1989	

NIFURTIMOX

TABLET; ORAL

LAMPIT

<u>+</u>	BAYER HEALTHCARE	<u>30MG</u>	<u>N213464</u>	<u>001</u>	Aug 06, 2020
<u>+</u> !		<u>120MG</u>	<u>N213464</u>	<u>002</u>	Aug 06, 2020

NILOTINIB HYDROCHLORIDE

CAPSULE; ORAL

TASIGNA

<u>+</u>	NOVARTIS	EQ 50MG BASE	<u>N022068</u>	<u>003</u>	Mar 22, 2018
<u>+</u>		EQ 150MG BASE	<u>N022068</u>	<u>002</u>	Jun 17, 2010
<u>+</u> !		EQ 200MG BASE	<u>N022068</u>	<u>001</u>	Oct 29, 2007

NILOTINIB TARTRATE

TABLET; ORAL

DANZITEN

<u>+</u>	AZURITY	EQ 71MG BASE	<u>N219293</u>	<u>001</u>	Nov 07, 2024
<u>+</u> !		EQ 95MG BASE	<u>N219293</u>	<u>002</u>	Nov 07, 2024

PRESCRIPTION DRUG PRODUCT LIST

NILUTAMIDE

TABLET; ORAL

NILANDRON

AB	+ !	ADVANZ PHARMA	150MG	N020169	002	Apr 30, 1999
-----------	------------	---------------	--------------	----------------	------------	--------------

NILUTAMIDE

AB		ANI PHARMS	150MG	A207631	001	Jul 15, 2016
-----------	--	------------	--------------	----------------	------------	--------------

NIMODIPINE

CAPSULE; ORAL

NIMODIPINE

AB	!	BIONPHARMA	30MG	A076740	001	Jan 17, 2008
-----------	----------	------------	-------------	----------------	------------	--------------

AB		HERITAGE	30MG	A077811	001	May 02, 2007
-----------	--	----------	-------------	----------------	------------	--------------

AB		THEPHARMANETWORK LLC	30MG	A090103	001	Apr 07, 2014
-----------	--	-------------------------	-------------	----------------	------------	--------------

SOLUTION; ORAL

NIMODIPINE

!		ANNORA PHARMA	3MG/ML	A216937	001	Jul 09, 2024
----------	--	---------------	--------	---------	-----	--------------

NYMALIZE

+ !		AZURITY	6MG/ML	N203340	002	Apr 08, 2020
------------	--	---------	--------	---------	-----	--------------

NINTEDANIB ESYLATE

CAPSULE; ORAL

OFEV

+		BOEHRINGER INGELHEIM	EQ 100MG BASE	N205832	001	Oct 15, 2014
----------	--	-------------------------	---------------	---------	-----	--------------

+ !			EQ 150MG BASE	N205832	002	Oct 15, 2014
------------	--	--	---------------	---------	-----	--------------

NIRAPARIB TOSYLATE

TABLET; ORAL

ZEJULA

+		GLAXOSMITHKLINE	EQ 100MG BASE	N214876	001	Apr 26, 2023
----------	--	-----------------	---------------	---------	-----	--------------

+			EQ 200MG BASE	N214876	002	Apr 26, 2023
----------	--	--	---------------	---------	-----	--------------

+ !			EQ 300MG BASE	N214876	003	Apr 26, 2023
------------	--	--	---------------	---------	-----	--------------

NIRMATRELVIR; RITONAVIR

TABLET; ORAL

PAXLOVID (COPACKAGED)

+ !		PFIZER	150MG;100MG	N217188	001	May 25, 2023
------------	--	--------	-------------	---------	-----	--------------

NIROGACESTAT HYDROBROMIDE

TABLET; ORAL

OGSIVEO

+		SPRINGWORKS	EQ 50MG BASE	N217677	001	Nov 27, 2023
----------	--	-------------	--------------	---------	-----	--------------

+			EQ 100MG BASE	N217677	002	Apr 04, 2024
----------	--	--	---------------	---------	-----	--------------

+ !			EQ 150MG BASE	N217677	003	Apr 04, 2024
------------	--	--	---------------	---------	-----	--------------

NISOLDIPINE

TABLET, EXTENDED RELEASE; ORAL

NISOLDIPINE

AB		MYLAN	8.5MG	A091001	001	Jan 26, 2011
-----------	--	-------	--------------	----------------	------------	--------------

AB			17MG	A091001	002	Jan 26, 2011
-----------	--	--	-------------	----------------	------------	--------------

AB			34MG	A091001	004	Jan 26, 2011
-----------	--	--	-------------	----------------	------------	--------------

SULAR

AB	+ !	COVIS	8.5MG	N020356	008	Jan 02, 2008
-----------	------------	-------	--------------	----------------	------------	--------------

AB	+ !		17MG	N020356	007	Jan 02, 2008
-----------	------------	--	-------------	----------------	------------	--------------

AB	+ !		34MG	N020356	005	Jan 02, 2008
-----------	------------	--	-------------	----------------	------------	--------------

NISOLDIPINE

!		MYLAN	20MG	A079051	001	Jul 25, 2008
----------	--	-------	------	---------	-----	--------------

			25.5MG	A091001	003	Jan 26, 2011
--	--	--	--------	---------	-----	--------------

			30MG	A079051	002	Jul 25, 2008
--	--	--	------	---------	-----	--------------

!			40MG	A079051	003	Jul 25, 2008
----------	--	--	------	---------	-----	--------------

NITAZOXANIDE

TABLET; ORAL

NITAZOXANIDE

!		RISING	500MG	A213820	001	Nov 27, 2020
----------	--	--------	-------	---------	-----	--------------

NITISINONE

CAPSULE; ORAL

NITISINONE

AB		ETON	2MG	A216201	001	May 25, 2023
-----------	--	------	------------	----------------	------------	--------------

AB			5MG	A216201	002	May 25, 2023
-----------	--	--	------------	----------------	------------	--------------

AB			10MG	A216201	003	May 25, 2023
-----------	--	--	-------------	----------------	------------	--------------

AB			20MG	A216201	004	May 25, 2023
-----------	--	--	-------------	----------------	------------	--------------

AB		MEDUNIK	2MG	A212390	001	May 26, 2022
-----------	--	---------	------------	----------------	------------	--------------

AB			5MG	A212390	002	May 26, 2022
-----------	--	--	------------	----------------	------------	--------------

AB			10MG	A212390	003	May 26, 2022
-----------	--	--	-------------	----------------	------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

NITISINONE

CAPSULE; ORAL

NITISINONE

<u>AB</u>		<u>20MG</u>	<u>A212390 004</u>	Apr 27, 2023
<u>AB</u>	NOVITIUM PHARMA	<u>2MG</u>	<u>A211041 001</u>	Aug 26, 2019
<u>AB</u>		<u>5MG</u>	<u>A211041 002</u>	Aug 26, 2019
<u>AB</u>		<u>10MG</u>	<u>A211041 003</u>	Aug 26, 2019

ORFADIN

<u>AB</u>	+	SWEDISH ORPHAN	<u>2MG</u>	<u>N021232 001</u>	Jan 18, 2002
<u>AB</u>	+		<u>5MG</u>	<u>N021232 002</u>	Jan 18, 2002
<u>AB</u>	+		<u>10MG</u>	<u>N021232 003</u>	Jan 18, 2002
<u>AB</u>	+	!	<u>20MG</u>	<u>N021232 004</u>	Jun 13, 2016

SUSPENSION; ORAL

ORFADIN

	+	SWEDISH ORPHAN	4MG/ML	N206356 001	Apr 22, 2016
--	---	----------------	--------	-------------	--------------

TABLET; ORAL

NITYR

	+	CYCLE	2MG	N209449 001	Jul 26, 2017
	+		5MG	N209449 002	Jul 26, 2017
	+	!	10MG	N209449 003	Jul 26, 2017

NITRIC OXIDE

GAS; INHALATION

INOMAX

<u>AA</u>	+	MALLINCKRODT HOSP	<u>800PPM</u>	<u>N020845 003</u>	Dec 23, 1999
-----------	---	-------------------	---------------	--------------------	--------------

NOXIVENT

<u>AA</u>		LINDE GAS EQUIP	<u>800PPM</u>	<u>A207141 002</u>	Oct 02, 2018
-----------	--	-----------------	---------------	--------------------	--------------

ULSPIRA

<u>AA</u>		AIRGAS THERAP	<u>800PPM</u>	<u>A203144 001</u>	Jul 27, 2023
-----------	--	---------------	---------------	--------------------	--------------

GENOSYL

	+	VERO BIOTECH INC	800PPM	N202860 001	Dec 20, 2019
--	---	------------------	--------	-------------	--------------

INOMAX

	+	MALLINCKRODT HOSP	4880PPM	N020845 004	Jan 17, 2023
--	---	-------------------	---------	-------------	--------------

NOXIVENT

	!	LINDE GAS EQUIP	100PPM	A207141 001	Oct 02, 2018
--	---	-----------------	--------	-------------	--------------

NITROFURANTOIN

SUSPENSION; ORAL

FURADANTIN

<u>AB</u>	+	CASPER PHARMA LLC	<u>25MG/5ML</u>	<u>N009175 001</u>	
-----------	---	-------------------	-----------------	--------------------	--

NITROFURANTOIN

<u>AB</u>		ACTAVIS MID ATLANTIC	<u>25MG/5ML</u>	<u>A205180 001</u>	May 03, 2016
<u>AB</u>		AMNEAL PHARMS	<u>25MG/5ML</u>	<u>A201679 001</u>	May 11, 2011
<u>AB</u>		AUROBINDO PHARMA	<u>25MG/5ML</u>	<u>A212607 001</u>	May 11, 2023
<u>AB</u>		BIOCON PHARMA	<u>25MG/5ML</u>	<u>A218346 001</u>	Oct 28, 2024
<u>AB</u>		NOSTRUM LABS INC	<u>25MG/5ML</u>	<u>A201355 001</u>	Aug 14, 2013
<u>AB</u>		NOVEL LABS INC	<u>25MG/5ML</u>	<u>A201693 001</u>	Sep 08, 2014
<u>AB</u>		NOVITIUM PHARMA	<u>25MG/5ML</u>	<u>A216385 001</u>	Apr 14, 2023

FURADANTIN

	+	CASPER PHARMA LLC	50MG/5ML	N009175 002	Jun 09, 2023
--	---	-------------------	----------	-------------	--------------

NITROFURANTOIN, MACROCRYSTALLINE

CAPSULE; ORAL

MACRODANTIN

<u>AB</u>	+	ALMATICA	<u>25MG</u>	<u>N016620 003</u>	
<u>AB</u>	+		<u>50MG</u>	<u>N016620 001</u>	
<u>AB</u>	+	!	<u>100MG</u>	<u>N016620 002</u>	

NITROFURANTOIN

<u>AB</u>		ALEMBIC	<u>25MG</u>	<u>A211935 001</u>	Jun 25, 2021
<u>AB</u>			<u>50MG</u>	<u>A211935 002</u>	Jun 25, 2021
<u>AB</u>			<u>100MG</u>	<u>A211935 003</u>	Jun 25, 2021
<u>AB</u>		AUROBINDO PHARMA USA	<u>25MG</u>	<u>A077025 002</u>	Oct 02, 2024
<u>AB</u>			<u>50MG</u>	<u>A074967 001</u>	Jul 09, 1997
<u>AB</u>			<u>100MG</u>	<u>A074967 002</u>	Jul 09, 1997
<u>AB</u>			<u>100MG</u>	<u>A077025 001</u>	Aug 18, 2004
<u>AB</u>		IMPAX LABS INC	<u>50MG</u>	<u>A073671 001</u>	Jan 28, 1993
<u>AB</u>			<u>100MG</u>	<u>A073652 001</u>	Jan 28, 1993
<u>AB</u>		MANKIND PHARMA	<u>25MG</u>	<u>A217272 001</u>	Mar 21, 2023
<u>AB</u>			<u>50MG</u>	<u>A217272 002</u>	Mar 21, 2023
<u>AB</u>			<u>100MG</u>	<u>A217272 003</u>	Mar 21, 2023
<u>AB</u>		NOVEL LABS INC	<u>50MG</u>	<u>A203233 001</u>	Jul 09, 2018
<u>AB</u>			<u>100MG</u>	<u>A203233 002</u>	Jul 09, 2018

PRESCRIPTION DRUG PRODUCT LIST

NITROFURANTOIN, MACROCRYSTALLINE

CAPSULE; ORAL

NITROFURANTOIN

<u>AB</u>	SUN PHARM INDUSTRIES	<u>25MG</u>	<u>A201722 001</u>	Feb 16, 2016
<u>AB</u>		<u>50MG</u>	<u>A201722 002</u>	Feb 16, 2016
<u>AB</u>		<u>100MG</u>	<u>A201722 003</u>	Feb 16, 2016
<u>AB</u>	ZYDUS PHARMS	<u>50MG</u>	<u>A205005 001</u>	Dec 12, 2017
<u>AB</u>		<u>100MG</u>	<u>A205005 002</u>	Dec 12, 2017

NITROFURANTOIN; NITROFURANTOIN, MACROCRYSTALLINE

CAPSULE; ORAL

MACROBID

<u>AB</u>	+! ALMATICA	<u>75MG; 25MG</u>	<u>N020064 001</u>	Dec 24, 1991
-----------	-------------	-------------------	--------------------	--------------

NITROFURANTOIN (MONOHYDRATE/MACROCRYSTALS)

<u>AB</u>	AMNEAL PHARMS	<u>75MG; 25MG</u>	<u>A207372 001</u>	May 15, 2017
<u>AB</u>	AUROBINDO PHARMA	<u>75MG; 25MG</u>	<u>A209225 001</u>	Mar 30, 2023
<u>AB</u>	INVENTIA	<u>75MG; 25MG</u>	<u>A211013 001</u>	Feb 18, 2022
<u>AB</u>	MANKIND PHARMA	<u>75MG; 25MG</u>	<u>A217357 001</u>	Jul 11, 2023
<u>AB</u>	OMSAV PHARMA	<u>75MG; 25MG</u>	<u>A217073 001</u>	Nov 03, 2023
<u>AB</u>	SANDOZ	<u>75MG; 25MG</u>	<u>A077066 001</u>	Apr 05, 2005
<u>AB</u>	SUNNY	<u>75MG; 25MG</u>	<u>A208516 001</u>	May 24, 2018
<u>AB</u>	WATSON LABS INC	<u>75MG; 25MG</u>	<u>A202250 001</u>	Jul 08, 2015

NITROGLYCERIN

AEROSOL, METERED; SUBLINGUAL

NITROMIST

+!	EVUS	0.4MG/SPRAY	N021780 001	Nov 02, 2006
----	------	-------------	-------------	--------------

FILM, EXTENDED RELEASE; TRANSDERMAL

NITROGLYCERIN

<u>AB2</u>	! MYLAN TECHNOLOGIES	<u>0.1MG/HR</u>	<u>A074559 004</u>	Feb 06, 1998
<u>AB2</u>	!	<u>0.2MG/HR</u>	<u>A074559 003</u>	Aug 30, 1996
<u>AB2</u>	!	<u>0.4MG/HR</u>	<u>A074559 002</u>	Aug 30, 1996
<u>AB2</u>	!	<u>0.6MG/HR</u>	<u>A074559 001</u>	Aug 30, 1996
<u>AB2</u>	ZYDUS PHARMS	<u>0.1MG/HR</u>	<u>A089885 002</u>	Oct 30, 2017
<u>AB2</u>		<u>0.2MG/HR</u>	<u>A089884 001</u>	Oct 30, 1998
<u>AB2</u>		<u>0.4MG/HR</u>	<u>A089885 001</u>	Oct 30, 1998
<u>AB2</u>		<u>0.6MG/HR</u>	<u>A089886 001</u>	Oct 30, 1998

NITRO-DUR

+!	USPHARMA	0.1MG/HR	N020145 001	Apr 04, 1995
+!		0.2MG/HR	N020145 002	Apr 04, 1995
+!		0.3MG/HR	N020145 003	Apr 04, 1995
+!		0.4MG/HR	N020145 004	Apr 04, 1995
+!		0.6MG/HR	N020145 005	Apr 04, 1995
+!		0.8MG/HR	N020145 006	Apr 04, 1995

INJECTABLE; INJECTION

NITROGLYCERIN

!	AM REGENT	5MG/ML	A072034 001	May 24, 1988
NITROGLYCERIN IN DEXTROSE 5%				
+!	BAXTER HLTHCARE	10MG/100ML	N019970 001	Dec 29, 1989
+!		20MG/100ML	N019970 002	Dec 29, 1989
+!		40MG/100ML	N019970 003	Dec 29, 1989

OINTMENT; INTRA-ANAL

NITROGLYCERIN

<u>AB</u>	ACRUX DDS	<u>0.4%</u>	<u>A212222 001</u>	Nov 29, 2024
<u>AB</u>	COSETTE	<u>0.4%</u>	<u>A216103 001</u>	Feb 16, 2024
<u>AB</u>	ENCUBE	<u>0.4%</u>	<u>A216452 001</u>	Dec 16, 2024

RECTIV

<u>AB</u>	+! ABEVIE	<u>0.4%</u>	<u>N021359 001</u>	Jun 21, 2011
-----------	-----------	-------------	--------------------	--------------

OINTMENT; TRANSDERMAL

NITROGLYCERIN

+!	FOUGERA PHARMS INC	2%	A087355 001	Jul 08, 1988
----	--------------------	----	-------------	--------------

SPRAY, METERED; SUBLINGUAL

NITROGLYCERIN

<u>AB</u>	PADAGIS ISRAEL	<u>0.4MG/SPRAY</u>	<u>A091496 001</u>	Sep 20, 2013
-----------	----------------	--------------------	--------------------	--------------

NITROLINGUAL PUMPSPRAY

<u>AB</u>	+! POHL BOSKAMP	<u>0.4MG/SPRAY</u>	<u>N018705 002</u>	Jan 10, 1997
-----------	-----------------	--------------------	--------------------	--------------

TABLET; SUBLINGUAL

NITROGLYCERIN

<u>AB</u>	AUROBINDO PHARMA	<u>0.3MG</u>	<u>A217879 001</u>	Oct 26, 2023
<u>AB</u>		<u>0.4MG</u>	<u>A217879 002</u>	Oct 26, 2023
<u>AB</u>		<u>0.6MG</u>	<u>A217879 003</u>	Oct 26, 2023
<u>AB</u>	DR REDDYS	<u>0.3MG</u>	<u>A208191 001</u>	Aug 26, 2016
<u>AB</u>		<u>0.4MG</u>	<u>A208191 002</u>	Aug 26, 2016

PRESCRIPTION DRUG PRODUCT LIST

NITROGLYCERIN

TABLET;SUBLINGUAL

NITROGLYCERIN

<u>AB</u>		<u>0.6MG</u>	<u>A208191</u>	<u>003</u>	Aug 26, 2016
<u>AB</u>	GLENMARK SPECLT	<u>0.3MG</u>	<u>A206391</u>	<u>001</u>	Sep 19, 2017
<u>AB</u>		<u>0.4MG</u>	<u>A206391</u>	<u>002</u>	Sep 19, 2017
<u>AB</u>		<u>0.6MG</u>	<u>A206391</u>	<u>003</u>	Sep 19, 2017
<u>AB</u>	MANKIND PHARMA	<u>0.3MG</u>	<u>A217970</u>	<u>001</u>	Dec 05, 2023
<u>AB</u>		<u>0.4MG</u>	<u>A217970</u>	<u>002</u>	Dec 05, 2023
<u>AB</u>		<u>0.6MG</u>	<u>A217970</u>	<u>003</u>	Dec 05, 2023
<u>AB</u>	NATCO	<u>0.3MG</u>	<u>A211604</u>	<u>001</u>	Apr 30, 2019
<u>AB</u>		<u>0.4MG</u>	<u>A211604</u>	<u>002</u>	Apr 30, 2019
<u>AB</u>		<u>0.6MG</u>	<u>A211604</u>	<u>003</u>	Apr 30, 2019
<u>AB</u>	RUBICON	<u>0.3MG</u>	<u>A209779</u>	<u>001</u>	May 03, 2021
<u>AB</u>		<u>0.4MG</u>	<u>A209779</u>	<u>002</u>	May 03, 2021
<u>AB</u>		<u>0.6MG</u>	<u>A209779</u>	<u>003</u>	May 03, 2021
<u>AB</u>	VIWIT PHARM	<u>0.3MG</u>	<u>A218583</u>	<u>001</u>	Oct 15, 2024
<u>AB</u>		<u>0.4MG</u>	<u>A218583</u>	<u>002</u>	Oct 15, 2024
<u>AB</u>		<u>0.6MG</u>	<u>A218583</u>	<u>003</u>	Oct 15, 2024
<u>NITROSTAT</u>					
<u>AB</u>	+ VIATRIS	<u>0.3MG</u>	<u>N021134</u>	<u>001</u>	May 01, 2000
<u>AB</u>	+	<u>0.4MG</u>	<u>N021134</u>	<u>002</u>	May 01, 2000
<u>AB</u>	+!	<u>0.6MG</u>	<u>N021134</u>	<u>003</u>	May 01, 2000

NIZATIDINE

CAPSULE;ORAL

NIZATIDINE

<u>AB</u>	DR REDDYS LABS LTD	<u>150MG</u>	<u>A077314</u>	<u>001</u>	Sep 15, 2005
<u>AB</u>		<u>300MG</u>	<u>A077314</u>	<u>002</u>	Sep 15, 2005
<u>AB</u>	EPIC PHARMA LLC	<u>150MG</u>	<u>A076178</u>	<u>001</u>	Jul 05, 2002
<u>AB</u>		<u>300MG</u>	<u>A076178</u>	<u>002</u>	Jul 05, 2002
<u>AB</u>	GLENMARK PHARMS INC	<u>150MG</u>	<u>A090618</u>	<u>001</u>	Jul 15, 2011
<u>AB</u>		<u>300MG</u>	<u>A090618</u>	<u>002</u>	Jul 15, 2011
<u>AB</u>	WATSON LABS	<u>150MG</u>	<u>A075616</u>	<u>001</u>	Jul 09, 2002
<u>AB</u>	!	<u>300MG</u>	<u>A075616</u>	<u>002</u>	Jul 09, 2002

NOREPINEPHRINE BITARTRATE

INJECTABLE;INJECTION

LEVOPHED

<u>AP</u>	+! HOSPIRA	<u>EQ 1MG BASE/ML</u>	<u>N007513</u>	<u>001</u>	
<u>NOREPINEPHRINE BITARTRATE</u>					
<u>AP</u>	AMNEAL	<u>EQ 1MG BASE/ML</u>	<u>A210839</u>	<u>001</u>	Dec 17, 2018
<u>AP</u>	BAXTER HLTHCARE CORP	<u>EQ 1MG BASE/ML</u>	<u>A040859</u>	<u>001</u>	Mar 27, 2012
<u>AP</u>	BRECKENRIDGE	<u>EQ 1MG BASE/ML</u>	<u>A214455</u>	<u>001</u>	Jan 22, 2021
<u>AP</u>	CAPLIN	<u>EQ 1MG BASE/ML</u>	<u>A217575</u>	<u>001</u>	Sep 15, 2023
<u>AP</u>	FRESENIUS KABI USA	<u>EQ 1MG BASE/ML</u>	<u>A211382</u>	<u>001</u>	Nov 03, 2020
<u>AP</u>	GLAND PHARMA LTD	<u>EQ 1MG BASE/ML</u>	<u>A214323</u>	<u>001</u>	May 06, 2021
<u>AP</u>	HIKMA	<u>EQ 1MG BASE/ML</u>	<u>A040462</u>	<u>001</u>	Oct 31, 2003
<u>AP</u>		<u>EQ 1MG BASE/ML</u>	<u>A203662</u>	<u>001</u>	Nov 07, 2018
<u>AP</u>	MEITHEAL	<u>EQ 1MG BASE/ML</u>	<u>A040455</u>	<u>001</u>	Mar 03, 2003
<u>AP</u>	MYLAN LABS LTD	<u>EQ 1MG BASE/ML</u>	<u>A211242</u>	<u>001</u>	Oct 04, 2018
<u>AP</u>	RISING	<u>EQ 1MG BASE/ML</u>	<u>A218758</u>	<u>001</u>	May 13, 2024
<u>AP</u>	SANDOZ	<u>EQ 1MG BASE/ML</u>	<u>A211359</u>	<u>001</u>	Oct 18, 2018

SOLUTION;INTRAVENOUS

NOREPINEPHRINE BITARTRATE IN 0.9% SODIUM CHLORIDE

+!	INFORLIFE	<u>EQ 4MG BASE/250ML (EQ 16MCG BASE/ML)</u>	<u>N215700</u>	<u>001</u>	Sep 15, 2022
+!		<u>EQ 8MG BASE/250ML (EQ 32MCG BASE/ML)</u>	<u>N215700</u>	<u>002</u>	Sep 15, 2022
+!		<u>EQ 16MG BASE/250ML (EQ 64MCG BASE/ML)</u>	<u>N215700</u>	<u>003</u>	Sep 15, 2022
+!	LONG GROVE PHARMS	<u>EQ 4MG BASE/250 ML (EQ 16MCG BASE/ML)</u>	<u>N214628</u>	<u>001</u>	Oct 06, 2022
+!		<u>EQ 8MG BASE/250ML (EQ 32MCG BASE/ML)</u>	<u>N214628</u>	<u>002</u>	Oct 06, 2022
+!		<u>EQ 16MG BASE/250ML (EQ 64MCG BASE/ML)</u>	<u>N214628</u>	<u>003</u>	Oct 06, 2022

NOREPINEPHRINE BITARTRATE IN 5% DEXTROSE

+!	BAXTER HLTHCARE CORP	<u>EQ 4MG BASE/250ML (EQ 16MCG BASE/ML)</u>	<u>N214313</u>	<u>001</u>	Jan 15, 2021
+!		<u>EQ 8MG BASE/250ML (EQ 32MCG BASE/ML)</u>	<u>N214313</u>	<u>002</u>	Jan 15, 2021
+!		<u>EQ 16MG BASE/250ML (EQ 64MCG BASE/ML)</u>	<u>N214313</u>	<u>003</u>	Nov 21, 2023

PRESCRIPTION DRUG PRODUCT LIST

NORETHINDRONE

TABLET; ORAL-28

CAMILA**AB1** DR REDDYS LABS SA **0.35MG** **A076177 001** Oct 21, 2002HEATHER**AB1** GLENMARK PHARMS LTD **0.35MG** **A090454 001** Apr 23, 2010INCASSIA**AB1** AUROBINDO PHARMA **0.35MG** **A207304 001** Sep 23, 2016NOR-QD**AB1** +! TEVA BRANDED PHARM **0.35MG** **N017060 001**NORETHINDRONE**AB1** LUPIN LTD **0.35MG** **A091325 001** Sep 19, 2011**AB1** NAARI PTE LTD **0.35MG** **A206807 001** Dec 13, 2016**AB1** NOVAST LABS **0.35MG** **A202014 001** Sep 13, 2013**AB1** XIROMED **0.35MG** **A201483 001** Jun 24, 2013EMZAHH**AB2** AUROBINDO PHARMA **0.35MG** **A216796 001** Jan 06, 2023ERRIN**AB2** DR REDDYS LABS SA **0.35MG** **A076225 001** Oct 21, 2002JENCYCLA**AB2** LUPIN LTD **0.35MG** **A091323 001** Mar 28, 2013NORETHINDRONE**AB2** GLENMARK PHARMS LTD **0.35MG** **A091209 001** Jul 22, 2010**AB2** NOVAST LABS **0.35MG** **A200961 001** Sep 13, 2013**AB2** ! XIROMED **0.35MG** **A200980 001** Jun 12, 2013NORETHINDRONE ACETATE

TABLET; ORAL

NORETHINDRONE ACETATE**AB** AMNEAL PHARMS **5MG** **A200275 001** Jul 30, 2012**AB** ! BARR **5MG** **A075951 001** May 25, 2001**AB** GLENMARK PHARMS LTD **5MG** **A091090 001** Jul 21, 2010**AB** NOVAST LABS **5MG** **A206490 001** Nov 05, 2018**AB** XIROMED **5MG** **A205278 001** Nov 10, 2016NORTRIPTYLINE HYDROCHLORIDE

CAPSULE; ORAL

NORTRIPTYLINE HYDROCHLORIDE**AB** DR REDDYS LABS SA **EQ 10MG BASE** **A073556 002** Mar 30, 1992**AB** **EQ 25MG BASE** **A073556 003** Mar 30, 1992**AB** **EQ 50MG BASE** **A073556 004** Mar 30, 1992**AB** **EQ 75MG BASE** **A073556 001** Mar 30, 1992**AB** TARO **EQ 10MG BASE** **A075520 004** May 08, 2000**AB** **EQ 25MG BASE** **A075520 003** May 08, 2000**AB** **EQ 50MG BASE** **A075520 001** May 08, 2000**AB** **EQ 75MG BASE** **A075520 002** May 08, 2000**AB** TEVA **EQ 10MG BASE** **A074132 001** Mar 27, 1995**AB** **EQ 25MG BASE** **A074132 002** Mar 27, 1995**AB** **EQ 50MG BASE** **A074132 003** Mar 27, 1995**AB** ! **EQ 75MG BASE** **A074132 004** Mar 27, 1995

SOLUTION; ORAL

NORTRIPTYLINE HYDROCHLORIDE**AA** ! PHARM ASSOC **EQ 10MG BASE/5ML** **A075606 001** Aug 23, 2000**AA** RUBICON **EQ 10MG BASE/5ML** **A217731 001** Aug 15, 2023NUSINERSEN SODIUM

SOLUTION; INTRATHECAL

SPINRAZA+! BIOGEN IDEC **EQ 12MG BASE/5ML (EQ 2.4MG BASE/ML)** **N209531 001** Dec 23, 2016NYSTATIN

CREAM; TOPICAL

NYSTATIN**AT** ACTAVIS MID **100,000 UNITS/GM** **A062949 001** Jun 13, 1988**AT** ATLANTIC **100,000 UNITS/GM** **A061966 001****AT** COSETTE **100,000 UNITS/GM** **A207733 001** Sep 26, 2017**AT** CROWN LABS INC **100,000 UNITS/GM** **A062129 001****AT** FOUGERA PHARMS **100,000 UNITS/GM** **A213566 001** Aug 10, 2021**AT** MACLEODS PHARMS LTD **100,000 UNITS/GM** **A062225 001****AT** PADAGIS US **100,000 UNITS/GM** **A064022 001** Jan 29, 1993**AT** ! TARO **100,000 UNITS/GM** **A212557 001** Jul 24, 2019**AT** TORRENT **100,000 UNITS/GM** **A212557 001**

PRESCRIPTION DRUG PRODUCT LIST

NYSTATIN

OINTMENT; TOPICAL

NYSTATIN

<u>AT</u>	ACTAVIS MID ATLANTIC	<u>100,000 UNITS/GM</u>	<u>A062840 001</u>	Nov 13, 1987
<u>AT</u>	COSETTE	<u>100,000 UNITS/GM</u>	<u>A209114 001</u>	Oct 06, 2017
<u>AT</u>	! FOUGERA PHARMS	<u>100,000 UNITS/GM</u>	<u>A062124 002</u>	Sep 23, 1982
<u>AT</u>	LYNE	<u>100,000 UNITS/GM</u>	<u>A209082 001</u>	May 21, 2018
<u>AT</u>	MACLEODS PHARMS LTD	<u>100,000 UNITS/GM</u>	<u>A213826 001</u>	Jan 14, 2021
<u>AT</u>	PADAGIS US	<u>100,000 UNITS/GM</u>	<u>A062472 001</u>	Feb 13, 1984
<u>AT</u>	TORRENT	<u>100,000 UNITS/GM</u>	<u>A211838 001</u>	Jan 28, 2019
<u>AT</u>	ZYDUS LIFESCIENCES	<u>100,000 UNITS/GM</u>	<u>A207767 001</u>	May 25, 2018

POWDER; TOPICAL

NYSTATIN

<u>AT</u>	ADRASTEIA PHARMA	<u>100,000 UNITS/GM</u>	<u>A065175 001</u>	Dec 17, 2004
<u>AT</u>	! DR REDDYS LABS SA	<u>100,000 UNITS/GM</u>	<u>A065203 001</u>	Jul 15, 2004
<u>AT</u>	EPIC PHARMA LLC	<u>100,000 UNITS/GM</u>	<u>A210532 001</u>	Apr 30, 2018
<u>AT</u>	LUPIN	<u>100,000 UNITS/GM</u>	<u>A065138 001</u>	Jul 23, 2004
<u>AT</u>	LYNE	<u>100,000 UNITS/GM</u>	<u>A208838 001</u>	May 30, 2017
<u>AT</u>	UPSHER SMITH LABS	<u>100,000 UNITS/GM</u>	<u>A065183 001</u>	May 03, 2005
<u>AT</u>	ZYDUS PHARMS	<u>100,000 UNITS/GM</u>	<u>A208581 001</u>	Jun 08, 2017

NYSTOP

<u>AT</u>	PADAGIS US	<u>100,000 UNITS/GM</u>	<u>A064118 001</u>	Aug 16, 1996
-----------	------------	-------------------------	--------------------	--------------

SUSPENSION; ORAL

NYSTATIN

<u>AA</u>	FOUGERA PHARMS INC	<u>100,000 UNITS/ML</u>	<u>A062517 001</u>	Jun 07, 1984
<u>AA</u>	GENUS	<u>100,000 UNITS/ML</u>	<u>A065148 001</u>	Jun 28, 2005
<u>AA</u>	! MEDLEY PHARMS	<u>100,000 UNITS/ML</u>	<u>A214346 001</u>	Mar 10, 2022
<u>AA</u>	PHARM ASSOC	<u>100,000 UNITS/ML</u>	<u>A203621 001</u>	Jan 07, 2016
<u>AA</u>	TARO	<u>100,000 UNITS/ML</u>	<u>A062876 001</u>	Feb 29, 1988
<u>AA</u>	WOCKHARDT BIO AG	<u>100,000 UNITS/ML</u>	<u>A062512 001</u>	Oct 29, 1984

TABLET; ORAL

NYSTATIN

<u>AA</u>	HERITAGE	<u>500,000 UNITS</u>	<u>A062474 001</u>	Dec 22, 1983
<u>AA</u>	! TEVA	<u>500,000 UNITS</u>	<u>A062506 001</u>	Jan 16, 1984

NYSTATIN; TRIAMCINOLONE ACETONIDE

CREAM; TOPICAL

NYSTATIN AND TRIAMCINOLONE ACETONIDE

<u>AT</u>	ALEMBIC	<u>100,000 UNITS/GM; 0.1%</u>	<u>A214090 001</u>	Mar 31, 2021
<u>AT</u>	AMNEAL	<u>100,000 UNITS/GM; 0.1%</u>	<u>A209990 001</u>	Feb 15, 2018
<u>AT</u>	CROWN LABS INC	<u>100,000 UNITS/GM; 0.1%</u>	<u>A207730 001</u>	Dec 26, 2017
<u>AT</u>	DR REDDYS	<u>100,000 UNITS/GM; 0.1%</u>	<u>A208326 001</u>	Oct 26, 2016
<u>AT</u>	FOUGERA PHARMS INC	<u>100,000 UNITS/GM; 0.1%</u>	<u>A062599 001</u>	Oct 08, 1985
<u>AT</u>	GLENMARK PHARMS LTD	<u>100,000 UNITS/GM; 0.1%</u>	<u>A208136 001</u>	Oct 24, 2016
<u>AT</u>	LUPIN LTD	<u>100,000 UNITS/GM; 0.1%</u>	<u>A208205 001</u>	May 31, 2018
<u>AT</u>	MACLEODS PHARMS LTD	<u>100,000 UNITS/GM; 0.1%</u>	<u>A214181 001</u>	Jul 13, 2022
<u>AT</u>	PADAGIS ISRAEL	<u>100,000 UNITS/GM; 0.1%</u>	<u>A208479 001</u>	Aug 14, 2017
<u>AT</u>	! TARO	<u>100,000 UNITS/GM; 0.1%</u>	<u>A062364 001</u>	Dec 22, 1987

OINTMENT; TOPICAL

NYSTATIN AND TRIAMCINOLONE ACETONIDE

<u>AT</u>	ALEMBIC	<u>100,000 UNITS/GM; 0.1%</u>	<u>A214186 001</u>	Mar 04, 2022
<u>AT</u>	DR REDDYS	<u>100,000 UNITS/GM; 0.1%</u>	<u>A207741 001</u>	Jan 31, 2017
<u>AT</u>	EPIC PHARMA LLC	<u>100,000 UNITS/GM; 0.1%</u>	<u>A207217 001</u>	Aug 04, 2017
<u>AT</u>	FOUGERA PHARMS INC	<u>100,000 UNITS/GM; 0.1%</u>	<u>A062602 001</u>	Oct 09, 1985
<u>AT</u>	GLENMARK PHARMS LTD	<u>100,000 UNITS/GM; 0.1%</u>	<u>A208300 001</u>	Jun 23, 2016
<u>AT</u>	MACLEODS PHARMS LTD	<u>100,000 UNITS/GM; 0.1%</u>	<u>A214751 001</u>	Aug 17, 2021
<u>AT</u>	PADAGIS ISRAEL	<u>100,000 UNITS/GM; 0.1%</u>	<u>A207380 001</u>	Dec 20, 2016
<u>AT</u>	RISING	<u>100,000 UNITS/GM; 0.1%</u>	<u>A206785 001</u>	Dec 29, 2016
<u>AT</u>	STRIDES PHARMA	<u>100,000 UNITS/GM; 0.1%</u>	<u>A210077 001</u>	Jan 29, 2018
<u>AT</u>	! TARO	<u>100,000 UNITS/GM; 0.1%</u>	<u>A063305 001</u>	Mar 29, 1993
<u>AT</u>	ZYDUS LIFESCIENCES	<u>100,000 UNITS/GM; 0.1%</u>	<u>A207764 001</u>	Nov 08, 2018

OBETICHOLIC ACID

TABLET; ORAL

OCALIVA

+	INTERCEPT PHARMS	5MG	N207999 001	May 27, 2016
	INC			
+	!	10MG	N207999 002	May 27, 2016

PRESCRIPTION DRUG PRODUCT LIST

OCTREOTIDE ACETATE

CAPSULE, DELAYED RELEASE;ORAL

MYCAPSSA

+! CHIESI

EQ 20MG BASE

N208232 001 Jun 26, 2020

INJECTABLE; INJECTION

OCTREOTIDE ACETATEAB TEVA PHARMS USA INCEQ 10MG BASE/VIALA210317 001 Dec 05, 2023ABEQ 20MG BASE/VIALA210317 002 Dec 05, 2023ABEQ 30MG BASE/VIALA210317 003 Dec 05, 2023SANDOSTATIN LARAB + NOVARTISEQ 10MG BASE/VIALN021008 001 Nov 25, 1998AB +EQ 20MG BASE/VIALN021008 002 Nov 25, 1998AB +!EQ 30MG BASE/VIALN021008 003 Nov 25, 1998OCTREOTIDE ACETATEAP FRESENIUS KABI USAEQ 0.2MG BASE/MLA077450 001 Feb 10, 2006APEQ 1MG BASE/MLA077450 002 Feb 10, 2006AP

GLAND PHARMA LTD

EQ 0.1MG BASE/MLA216839 002 Jun 22, 2023APEQ 0.2MG BASE/MLA216807 001 Jun 13, 2023APEQ 0.5MG BASE/MLA216839 001 Jun 22, 2023APEQ 1MG BASE/MLA216807 002 Jun 13, 2023AP

HERITAGE

EQ 0.05MG BASE/MLA204669 001 Dec 27, 2018APEQ 0.1MG BASE/MLA204669 002 Dec 27, 2018APEQ 0.2MG BASE/MLA203765 001 Sep 07, 2018APEQ 0.5MG BASE/MLA204669 003 Dec 27, 2018APEQ 1MG BASE/MLA203765 002 Sep 07, 2018AP

MEITHEAL

EQ 0.05MG BASE/MLA075957 001 Oct 03, 2005APEQ 0.1MG BASE/MLA075957 002 Oct 03, 2005APEQ 0.2MG BASE/MLA075959 001 Nov 21, 2005APEQ 0.5MG BASE/MLA075957 003 Oct 03, 2005APEQ 1MG BASE/MLA075959 002 Nov 21, 2005AP

SAGENT PHARMS INC

EQ 0.2MG BASE/MLA091041 001 Nov 12, 2013APEQ 1MG BASE/MLA091041 002 Nov 12, 2013AP !

WEST-WARD PHARMS

EQ 0.2MG BASE/MLA076330 001 Apr 08, 2005

INT

AP !EQ 1MG BASE/MLA076330 002 Apr 08, 2005OCTREOTIDE ACETATE (PRESERVATIVE FREE)AP FRESENIUS KABI USAEQ 0.05MG BASE/MLA077457 001 Feb 10, 2006APEQ 0.1MG BASE/MLA077457 002 Feb 10, 2006APEQ 0.5MG BASE/MLA077457 003 Feb 10, 2006AP

MYLAN INSTITUTIONAL

EQ 0.05MG BASE/MLA079198 001 Feb 10, 2011APEQ 0.1MG BASE/MLA079198 002 Feb 10, 2011APEQ 0.5MG BASE/MLA079198 003 Feb 10, 2011AP

SAGENT PHARMS INC

EQ 0.05MG BASE/MLA090834 001 Nov 12, 2013APEQ 0.1MG BASE/MLA090834 002 Nov 12, 2013APEQ 0.5MG BASE/MLA090834 003 Nov 12, 2013AP !

WEST-WARD PHARMS

EQ 0.05MG BASE/MLA076313 001 Mar 28, 2005

INT

AP !EQ 0.1MG BASE/MLA076313 003 Mar 28, 2005AP !EQ 0.5MG BASE/MLA076313 002 Mar 28, 2005SANDOSTATINAP +! NOVARTISEQ 0.05MG BASE/MLN019667 001 Oct 21, 1988AP +!EQ 0.1MG BASE/MLN019667 002 Oct 21, 1988AP +!EQ 0.5MG BASE/MLN019667 003 Oct 21, 1988

SOLUTION; SUBCUTANEOUS

BYNFEZIA PEN

+! SUN PHARM

EQ 7MG BASE/2.8ML (EQ 2.5MG BASE/ML)

N213224 001 Sep 27, 2024

ODEVIXIBAT

CAPSULE; ORAL

BYLVAY

+ IPSEN

0.4MG

N215498 002 Jul 20, 2021

+!

1.2MG

N215498 004 Jul 20, 2021

CAPSULE, PELLETS; ORAL

BYLVAY

+ IPSEN

0.2MG

N215498 001 Jul 20, 2021

+!

0.6MG

N215498 003 Jul 20, 2021

OFLOXACIN

SOLUTION/DROPS; OPHTHALMIC

OCUFLOXAT +! ALLERGAN0.3%N019921 001 Jul 30, 1993OFLOXACINAT ALTAIRE PHARMS INC0.3%A202692 001 Apr 29, 2013AT

AMNEAL

0.3%A211524 001 Mar 01, 2024

PRESCRIPTION DRUG PRODUCT LIST

OFLOXACIN

SOLUTION/DROPS;OPHTHALMIC

OFLOXACIN

AT	APOTEX INC	<u>0.3%</u>	<u>A076513</u>	<u>001</u>	May 14, 2004
AT	BAUSCH AND LOMB	<u>0.3%</u>	<u>A076622</u>	<u>001</u>	May 14, 2004
AT	CAPLIN	<u>0.3%</u>	<u>A217904</u>	<u>001</u>	Apr 12, 2024
AT	FDC LTD	<u>0.3%</u>	<u>A078559</u>	<u>001</u>	Feb 25, 2009
AT	MANKIND PHARMA	<u>0.3%</u>	<u>A215886</u>	<u>001</u>	Nov 23, 2022
AT	SENTISS	<u>0.3%</u>	<u>A076407</u>	<u>001</u>	Apr 15, 2008
AT	SOMERSET	<u>0.3%</u>	<u>A213597</u>	<u>001</u>	Aug 01, 2024

SOLUTION/DROPS;OTIC

OFLOXACIN

AT	AMNEAL	<u>0.3%</u>	<u>A211525</u>	<u>001</u>	Aug 30, 2019
AT	APOTEX INC	<u>0.3%</u>	<u>A076527</u>	<u>001</u>	Sep 28, 2007
AT	! BAUSCH AND LOMB	<u>0.3%</u>	<u>A076128</u>	<u>001</u>	Mar 17, 2008
AT	CAPLIN	<u>0.3%</u>	<u>A217903</u>	<u>001</u>	Jan 05, 2024
AT	MANKIND PHARMA	<u>0.3%</u>	<u>A216130</u>	<u>001</u>	Jul 14, 2022
AT	SOMERSET THERAPS LLC	<u>0.3%</u>	<u>A216328</u>	<u>001</u>	Oct 15, 2024

TABLET;ORAL

OFLOXACIN

AB	CADILA PHARMS LTD	<u>200MG</u>	<u>A091656</u>	<u>001</u>	Sep 18, 2014
AB		<u>300MG</u>	<u>A091656</u>	<u>002</u>	Sep 18, 2014
AB		<u>400MG</u>	<u>A091656</u>	<u>003</u>	Sep 18, 2014
AB	CHARTWELL RX	<u>400MG</u>	<u>A076093</u>	<u>003</u>	Sep 02, 2003
AB	DR REDDYS LABS LTD	<u>200MG</u>	<u>A077098</u>	<u>001</u>	Feb 10, 2006
AB		<u>300MG</u>	<u>A077098</u>	<u>002</u>	Feb 10, 2006
AB		<u>400MG</u>	<u>A077098</u>	<u>003</u>	Feb 10, 2006
AB	TEVA	<u>200MG</u>	<u>A076182</u>	<u>001</u>	Sep 02, 2003
AB		<u>300MG</u>	<u>A076182</u>	<u>002</u>	Sep 02, 2003
AB	!	<u>400MG</u>	<u>A076182</u>	<u>003</u>	Sep 02, 2003

OLANZAPINE

INJECTABLE;INTRAMUSCULAR

OLANZAPINE

AP	AM REGENT	<u>10MG/VIAL</u>	<u>A201741</u>	<u>001</u>	Mar 20, 2012
AP	ASPIRO	<u>10MG/VIAL</u>	<u>A217466</u>	<u>001</u>	Mar 22, 2023
AP	EUGIA PHARMA	<u>10MG/VIAL</u>	<u>A210968</u>	<u>001</u>	Oct 22, 2020
AP	SANDOZ INC	<u>10MG/VIAL</u>	<u>A201588</u>	<u>001</u>	Oct 24, 2011

ZYPREXA

AP	+! CHEPLAPHARM	<u>10MG/VIAL</u>	<u>N021253</u>	<u>001</u>	Mar 29, 2004
-----------	----------------	------------------	-----------------------	-------------------	--------------

TABLET;ORAL

OLANZAPINE

AB	ALKEM LABS LTD	<u>2.5MG</u>	<u>A202295</u>	<u>001</u>	Oct 20, 2015
AB		<u>5MG</u>	<u>A202295</u>	<u>002</u>	Oct 20, 2015
AB		<u>7.5MG</u>	<u>A202295</u>	<u>003</u>	Oct 20, 2015
AB		<u>10MG</u>	<u>A202295</u>	<u>004</u>	Oct 20, 2015
AB		<u>15MG</u>	<u>A202295</u>	<u>005</u>	Oct 20, 2015
AB		<u>20MG</u>	<u>A202295</u>	<u>006</u>	Oct 20, 2015
AB	APOTEX INC	<u>2.5MG</u>	<u>A090798</u>	<u>001</u>	Apr 23, 2012
AB		<u>5MG</u>	<u>A090798</u>	<u>002</u>	Apr 23, 2012
AB		<u>7.5MG</u>	<u>A090798</u>	<u>003</u>	Apr 23, 2012
AB		<u>10MG</u>	<u>A090798</u>	<u>004</u>	Apr 23, 2012
AB		<u>15MG</u>	<u>A090798</u>	<u>005</u>	Apr 23, 2012
AB		<u>20MG</u>	<u>A090798</u>	<u>006</u>	Apr 23, 2012
AB	AUROBINDO PHARMA LTD	<u>2.5MG</u>	<u>A202050</u>	<u>001</u>	Apr 23, 2012
AB		<u>5MG</u>	<u>A202050</u>	<u>002</u>	Apr 23, 2012
AB		<u>7.5MG</u>	<u>A202050</u>	<u>003</u>	Apr 23, 2012
AB		<u>10MG</u>	<u>A202050</u>	<u>004</u>	Apr 23, 2012
AB		<u>15MG</u>	<u>A202050</u>	<u>005</u>	Apr 23, 2012
AB		<u>20MG</u>	<u>A202050</u>	<u>006</u>	Apr 23, 2012
AB	CADILA PHARMS LTD	<u>2.5MG</u>	<u>A210022</u>	<u>001</u>	Feb 24, 2023
AB		<u>5MG</u>	<u>A210022</u>	<u>002</u>	Feb 24, 2023
AB		<u>7.5MG</u>	<u>A210022</u>	<u>003</u>	Feb 24, 2023
AB		<u>10MG</u>	<u>A210022</u>	<u>004</u>	Feb 24, 2023
AB		<u>15MG</u>	<u>A210022</u>	<u>005</u>	Feb 24, 2023
AB		<u>20MG</u>	<u>A210022</u>	<u>006</u>	Feb 24, 2023
AB	CHARTWELL MOLECULAR	<u>2.5MG</u>	<u>A203333</u>	<u>001</u>	Mar 15, 2016
AB		<u>5MG</u>	<u>A203333</u>	<u>002</u>	Mar 15, 2016
AB		<u>7.5MG</u>	<u>A203333</u>	<u>003</u>	Mar 15, 2016
AB		<u>10MG</u>	<u>A203333</u>	<u>004</u>	Mar 15, 2016
AB		<u>15MG</u>	<u>A203333</u>	<u>005</u>	Mar 15, 2016

PRESCRIPTION DRUG PRODUCT LIST

OLANZAPINE

TABLET; ORAL

OLANZAPINE

<u>AB</u>		<u>20MG</u>	<u>A203333</u>	<u>006</u>	Mar 15, 2016
<u>AB</u>	DR REDDYS LABS LTD	<u>2.5MG</u>	<u>A076255</u>	<u>001</u>	Apr 23, 2012
<u>AB</u>		<u>5MG</u>	<u>A076255</u>	<u>002</u>	Apr 23, 2012
<u>AB</u>		<u>7.5MG</u>	<u>A076255</u>	<u>003</u>	Apr 23, 2012
<u>AB</u>		<u>10MG</u>	<u>A076255</u>	<u>004</u>	Apr 23, 2012
<u>AB</u>		<u>15MG</u>	<u>A076133</u>	<u>001</u>	Apr 23, 2012
<u>AB</u>		<u>20MG</u>	<u>A076133</u>	<u>002</u>	Oct 24, 2011
<u>AB</u>	MACLEODS PHARMS LTD	<u>2.5MG</u>	<u>A202862</u>	<u>001</u>	Aug 15, 2014
<u>AB</u>		<u>5MG</u>	<u>A202862</u>	<u>002</u>	Aug 15, 2014
<u>AB</u>		<u>7.5MG</u>	<u>A202862</u>	<u>003</u>	Aug 15, 2014
<u>AB</u>		<u>10MG</u>	<u>A202862</u>	<u>004</u>	Aug 15, 2014
<u>AB</u>		<u>15MG</u>	<u>A202862</u>	<u>005</u>	Aug 15, 2014
<u>AB</u>		<u>20MG</u>	<u>A202862</u>	<u>006</u>	Aug 15, 2014
<u>AB</u>	ORBION PHARMS	<u>2.5MG</u>	<u>A202287</u>	<u>001</u>	Apr 23, 2012
<u>AB</u>		<u>5MG</u>	<u>A202287</u>	<u>002</u>	Apr 23, 2012
<u>AB</u>		<u>7.5MG</u>	<u>A202287</u>	<u>003</u>	Apr 23, 2012
<u>AB</u>		<u>10MG</u>	<u>A202287</u>	<u>004</u>	Apr 23, 2012
<u>AB</u>		<u>15MG</u>	<u>A202287</u>	<u>005</u>	Apr 23, 2012
<u>AB</u>		<u>20MG</u>	<u>A202287</u>	<u>006</u>	Apr 23, 2012
<u>AB</u>	QILU	<u>2.5MG</u>	<u>A204319</u>	<u>001</u>	Jan 27, 2016
<u>AB</u>		<u>5MG</u>	<u>A204319</u>	<u>002</u>	Jan 27, 2016
<u>AB</u>		<u>7.5MG</u>	<u>A204319</u>	<u>003</u>	Jan 27, 2016
<u>AB</u>		<u>10MG</u>	<u>A204319</u>	<u>004</u>	Jan 27, 2016
<u>AB</u>		<u>15MG</u>	<u>A204319</u>	<u>005</u>	Jan 27, 2016
<u>AB</u>		<u>20MG</u>	<u>A204319</u>	<u>006</u>	Jan 27, 2016
<u>AB</u>	ZYDUS PHARMS	<u>2.5MG</u>	<u>A090459</u>	<u>001</u>	Jul 16, 2018
<u>AB</u>		<u>5MG</u>	<u>A090459</u>	<u>002</u>	Jul 16, 2018
<u>AB</u>		<u>7.5MG</u>	<u>A090459</u>	<u>003</u>	Jul 16, 2018
<u>AB</u>		<u>10MG</u>	<u>A090459</u>	<u>004</u>	Jul 16, 2018
<u>AB</u>		<u>15MG</u>	<u>A090459</u>	<u>005</u>	Jul 16, 2018
<u>AB</u>		<u>20MG</u>	<u>A090459</u>	<u>006</u>	Jul 16, 2018

ZYPREXA

<u>AB</u>	+	CHEPLAPHARM	<u>2.5MG</u>	<u>N020592</u>	<u>001</u>	Sep 30, 1996
<u>AB</u>	+	!	<u>5MG</u>	<u>N020592</u>	<u>002</u>	Sep 30, 1996
<u>AB</u>	+		<u>7.5MG</u>	<u>N020592</u>	<u>003</u>	Sep 30, 1996
<u>AB</u>	+		<u>10MG</u>	<u>N020592</u>	<u>004</u>	Sep 30, 1996
<u>AB</u>	+		<u>15MG</u>	<u>N020592</u>	<u>005</u>	Sep 09, 1997
<u>AB</u>	+		<u>20MG</u>	<u>N020592</u>	<u>006</u>	Sep 09, 1997

TABLET, ORALLY DISINTEGRATING; ORAL

OLANZAPINE

<u>AB</u>	APOTEX INC	<u>5MG</u>	<u>A091265</u>	<u>001</u>	Oct 24, 2011
<u>AB</u>		<u>10MG</u>	<u>A091265</u>	<u>002</u>	Oct 24, 2011
<u>AB</u>		<u>15MG</u>	<u>A091265</u>	<u>003</u>	Oct 24, 2011
<u>AB</u>		<u>20MG</u>	<u>A091265</u>	<u>004</u>	Oct 24, 2011
<u>AB</u>	AUROBINDO PHARMA LTD	<u>5MG</u>	<u>A203708</u>	<u>001</u>	May 15, 2014
<u>AB</u>		<u>10MG</u>	<u>A203708</u>	<u>002</u>	May 15, 2014
<u>AB</u>		<u>15MG</u>	<u>A203708</u>	<u>003</u>	May 15, 2014
<u>AB</u>		<u>20MG</u>	<u>A203708</u>	<u>004</u>	May 15, 2014
<u>AB</u>	BARR LABS INC	<u>5MG</u>	<u>A077243</u>	<u>001</u>	Jan 30, 2012
<u>AB</u>		<u>10MG</u>	<u>A077243</u>	<u>002</u>	Jan 30, 2012
<u>AB</u>		<u>15MG</u>	<u>A077243</u>	<u>003</u>	Jan 30, 2012
<u>AB</u>		<u>20MG</u>	<u>A077243</u>	<u>004</u>	Jan 30, 2012
<u>AB</u>	CHARTWELL MOLECULAR	<u>5MG</u>	<u>A203456</u>	<u>001</u>	Mar 16, 2016
<u>AB</u>		<u>10MG</u>	<u>A203456</u>	<u>002</u>	Mar 16, 2016
<u>AB</u>		<u>15MG</u>	<u>A203456</u>	<u>003</u>	Mar 16, 2016
<u>AB</u>		<u>20MG</u>	<u>A203456</u>	<u>004</u>	Mar 16, 2016
<u>AB</u>	DR REDDYS LABS LTD	<u>5MG</u>	<u>A076534</u>	<u>001</u>	Oct 24, 2011
<u>AB</u>		<u>10MG</u>	<u>A076534</u>	<u>002</u>	Oct 24, 2011
<u>AB</u>		<u>15MG</u>	<u>A076534</u>	<u>003</u>	Oct 24, 2011
<u>AB</u>		<u>20MG</u>	<u>A076534</u>	<u>004</u>	Oct 24, 2011
<u>AB</u>	HEC PHARM	<u>5MG</u>	<u>A208146</u>	<u>001</u>	Jul 02, 2018
<u>AB</u>		<u>10MG</u>	<u>A208146</u>	<u>002</u>	Jul 02, 2018
<u>AB</u>		<u>15MG</u>	<u>A208146</u>	<u>003</u>	Jul 02, 2018
<u>AB</u>		<u>20MG</u>	<u>A208146</u>	<u>004</u>	Jul 02, 2018
<u>AB</u>	HISUN PHARM HANGZHOU	<u>5MG</u>	<u>A206892</u>	<u>001</u>	Dec 31, 2020
<u>AB</u>		<u>10MG</u>	<u>A206892</u>	<u>002</u>	Dec 31, 2020
<u>AB</u>		<u>15MG</u>	<u>A206892</u>	<u>003</u>	Dec 31, 2020
<u>AB</u>		<u>20MG</u>	<u>A206892</u>	<u>004</u>	Dec 31, 2020

PRESCRIPTION DRUG PRODUCT LIST

OLANZAPINE

TABLET, ORALLY DISINTEGRATING;ORAL

OLANZAPINE

<u>AB</u>	JUBILANT GENERICS	<u>5MG</u>	<u>A200221</u>	<u>001</u>	Sep 12, 2012
<u>AB</u>		<u>10MG</u>	<u>A200221</u>	<u>002</u>	Sep 12, 2012
<u>AB</u>		<u>15MG</u>	<u>A200221</u>	<u>003</u>	Sep 12, 2012
<u>AB</u>		<u>20MG</u>	<u>A200221</u>	<u>004</u>	Sep 12, 2012
<u>AB</u>	MACLEODS PHARMS LTD	<u>5MG</u>	<u>A203044</u>	<u>001</u>	Feb 20, 2015
<u>AB</u>		<u>10MG</u>	<u>A203044</u>	<u>002</u>	Feb 20, 2015
<u>AB</u>		<u>15MG</u>	<u>A203044</u>	<u>003</u>	Feb 20, 2015
<u>AB</u>		<u>20MG</u>	<u>A203044</u>	<u>004</u>	Feb 20, 2015
<u>AB</u>	MYLAN	<u>5MG</u>	<u>A202285</u>	<u>001</u>	May 12, 2014
<u>AB</u>		<u>10MG</u>	<u>A202285</u>	<u>002</u>	May 12, 2014
<u>AB</u>		<u>15MG</u>	<u>A202285</u>	<u>003</u>	May 12, 2014
<u>AB</u>		<u>20MG</u>	<u>A202285</u>	<u>004</u>	May 12, 2014
<u>AB</u>	ORBION PHARMS	<u>5MG</u>	<u>A202937</u>	<u>001</u>	Mar 02, 2015
<u>AB</u>		<u>10MG</u>	<u>A202937</u>	<u>002</u>	Mar 02, 2015
<u>AB</u>		<u>15MG</u>	<u>A202937</u>	<u>003</u>	Mar 02, 2015
<u>AB</u>		<u>20MG</u>	<u>A202937</u>	<u>004</u>	Mar 02, 2015
<u>AB</u>	STRIDES PHARMA	<u>5MG</u>	<u>A078109</u>	<u>001</u>	Oct 24, 2011
<u>AB</u>		<u>10MG</u>	<u>A078109</u>	<u>002</u>	Oct 24, 2011
<u>AB</u>		<u>15MG</u>	<u>A078109</u>	<u>003</u>	Oct 24, 2011
<u>AB</u>		<u>20MG</u>	<u>A078109</u>	<u>004</u>	Oct 24, 2011
<u>AB</u>	TORRENT	<u>5MG</u>	<u>A091415</u>	<u>001</u>	Oct 25, 2011
<u>AB</u>		<u>10MG</u>	<u>A091415</u>	<u>002</u>	Oct 25, 2011
<u>AB</u>		<u>15MG</u>	<u>A091415</u>	<u>003</u>	Oct 25, 2011
<u>AB</u>		<u>20MG</u>	<u>A091415</u>	<u>004</u>	Oct 25, 2011
<u>ZYPREXA ZYDIS</u>					
<u>AB</u>	+! CHEPLAPHARM	<u>5MG</u>	<u>N021086</u>	<u>001</u>	Apr 06, 2000
<u>AB</u>	+	<u>10MG</u>	<u>N021086</u>	<u>002</u>	Apr 06, 2000
<u>AB</u>	+	<u>15MG</u>	<u>N021086</u>	<u>003</u>	Apr 06, 2000
<u>AB</u>	+	<u>20MG</u>	<u>N021086</u>	<u>004</u>	Apr 06, 2000

OLANZAPINE PAMOATE

SUSPENSION, EXTENDED RELEASE;INTRAMUSCULAR

ZYPREXA RELPREVV

+	CHEPLAPHARM	EQ 210MG BASE/VIAL	N022173	001	Dec 11, 2009
+		EQ 300MG BASE/VIAL	N022173	002	Dec 11, 2009
+!		EQ 405MG BASE/VIAL	N022173	003	Dec 11, 2009

OLANZAPINE; SAMIDORPHAN L-MALATE

TABLET;ORAL

LYBALVI

+!	ALKERMES INC	5MG;EQ 10MG BASE	N213378	001	May 28, 2021
+		10MG;EQ 10MG BASE	N213378	002	May 28, 2021
+		15MG;EQ 10MG BASE	N213378	003	May 28, 2021
+		20MG;EQ 10MG BASE	N213378	004	May 28, 2021

OLAPARIB

TABLET;ORAL

LYNPARZA

+	ASTRAZENECA	100MG	N208558	001	Aug 17, 2017
+!		150MG	N208558	002	Aug 17, 2017

OLEZARSEN SODIUM

SOLUTION;SUBCUTANEOUS

TRYNGOLZA (AUTOINJECTOR)

+!	IONIS PHARMS INC	EQ 80MG BASE/0.8ML (EQ 80MG BASE/0.8ML)	N218614	001	Dec 19, 2024
----	------------------	---	---------	-----	--------------

OLICERIDINE

SOLUTION;INTRAVENOUS

OLINVYK

+!	TREVENA	1MG/ML (1MG/ML)	N210730	001	Oct 30, 2020
+!		2MG/2ML (1MG/ML)	N210730	002	Oct 30, 2020

OLIVE OIL; SOYBEAN OIL

EMULSION;INTRAVENOUS

CLINOLIPID 20%

+!	BAXTER HLTHCARE CORP	16%(160GM/1000ML);4% (40GM/1000ML)	N204508	001	Oct 03, 2013
----	----------------------	------------------------------------	---------	-----	--------------

PRESCRIPTION DRUG PRODUCT LIST

OLMESARTAN MEDOXOMIL

TABLET; ORAL

BENICAR

<u>AB</u>	+	COSETTE	<u>5MG</u>	<u>N021286</u>	<u>001</u>	Apr 25, 2002
<u>AB</u>	+		<u>20MG</u>	<u>N021286</u>	<u>003</u>	Apr 25, 2002
<u>AB</u>	+	!	<u>40MG</u>	<u>N021286</u>	<u>004</u>	Apr 25, 2002

OLMESARTAN MEDOXOMIL

<u>AB</u>		ACCORD HLTHCARE	<u>5MG</u>	<u>A207662</u>	<u>001</u>	Apr 24, 2017
<u>AB</u>			<u>20MG</u>	<u>A207662</u>	<u>002</u>	Apr 24, 2017
<u>AB</u>			<u>40MG</u>	<u>A207662</u>	<u>003</u>	Apr 24, 2017
<u>AB</u>		ALEMBIC	<u>5MG</u>	<u>A203012</u>	<u>001</u>	Apr 24, 2017
<u>AB</u>			<u>20MG</u>	<u>A203012</u>	<u>002</u>	Apr 24, 2017
<u>AB</u>			<u>40MG</u>	<u>A203012</u>	<u>003</u>	Apr 24, 2017
<u>AB</u>		ALKEM LABS LTD	<u>5MG</u>	<u>A206763</u>	<u>001</u>	Apr 24, 2017
<u>AB</u>			<u>20MG</u>	<u>A206763</u>	<u>002</u>	Apr 24, 2017
<u>AB</u>			<u>40MG</u>	<u>A206763</u>	<u>003</u>	Apr 24, 2017
<u>AB</u>		AUROBINDO PHARMA	<u>5MG</u>	<u>A204798</u>	<u>001</u>	Apr 24, 2017
<u>AB</u>			<u>20MG</u>	<u>A204798</u>	<u>002</u>	Apr 24, 2017
<u>AB</u>			<u>40MG</u>	<u>A204798</u>	<u>003</u>	Apr 24, 2017
<u>AB</u>		CADILA PHARMS LTD	<u>5MG</u>	<u>A210682</u>	<u>001</u>	Nov 08, 2024
<u>AB</u>			<u>20MG</u>	<u>A210682</u>	<u>002</u>	Nov 08, 2024
<u>AB</u>			<u>40MG</u>	<u>A210682</u>	<u>003</u>	Nov 08, 2024
<u>AB</u>		CHARTWELL RX	<u>5MG</u>	<u>A206227</u>	<u>001</u>	Jan 25, 2024
<u>AB</u>			<u>20MG</u>	<u>A206227</u>	<u>002</u>	Jan 25, 2024
<u>AB</u>			<u>40MG</u>	<u>A206227</u>	<u>003</u>	Jan 25, 2024
<u>AB</u>		GLENMARK PHARMS LTD	<u>5MG</u>	<u>A203281</u>	<u>001</u>	May 25, 2017
<u>AB</u>			<u>20MG</u>	<u>A203281</u>	<u>002</u>	May 25, 2017
<u>AB</u>			<u>40MG</u>	<u>A203281</u>	<u>003</u>	May 25, 2017
<u>AB</u>		HETERO LABS LTD V	<u>5MG</u>	<u>A205499</u>	<u>001</u>	Oct 31, 2024
<u>AB</u>			<u>20MG</u>	<u>A205499</u>	<u>002</u>	Oct 31, 2024
<u>AB</u>			<u>40MG</u>	<u>A205499</u>	<u>003</u>	Oct 31, 2024
<u>AB</u>		INVENTIA	<u>5MG</u>	<u>A208659</u>	<u>001</u>	May 18, 2020
<u>AB</u>			<u>20MG</u>	<u>A208659</u>	<u>002</u>	May 18, 2020
<u>AB</u>			<u>40MG</u>	<u>A208659</u>	<u>003</u>	May 18, 2020
<u>AB</u>		MACLEODS PHARMS LTD	<u>5MG</u>	<u>A204814</u>	<u>001</u>	Apr 24, 2017
<u>AB</u>			<u>20MG</u>	<u>A204814</u>	<u>002</u>	Apr 24, 2017
<u>AB</u>			<u>40MG</u>	<u>A204814</u>	<u>003</u>	Apr 24, 2017
<u>AB</u>		MICRO LABS	<u>5MG</u>	<u>A206372</u>	<u>001</u>	Sep 17, 2019
<u>AB</u>			<u>20MG</u>	<u>A206372</u>	<u>002</u>	Sep 17, 2019
<u>AB</u>			<u>40MG</u>	<u>A206372</u>	<u>003</u>	Sep 17, 2019
<u>AB</u>		MSN	<u>5MG</u>	<u>A217399</u>	<u>001</u>	Jan 18, 2023
<u>AB</u>			<u>20MG</u>	<u>A217399</u>	<u>002</u>	Jan 18, 2023
<u>AB</u>			<u>40MG</u>	<u>A217399</u>	<u>003</u>	Jan 18, 2023
<u>AB</u>		PRINSTON INC	<u>5MG</u>	<u>A206720</u>	<u>001</u>	Dec 02, 2022
<u>AB</u>			<u>20MG</u>	<u>A206720</u>	<u>002</u>	Dec 02, 2022
<u>AB</u>			<u>40MG</u>	<u>A206720</u>	<u>003</u>	Dec 02, 2022
<u>AB</u>		QILU	<u>5MG</u>	<u>A210552</u>	<u>001</u>	Jan 10, 2019
<u>AB</u>			<u>20MG</u>	<u>A210552</u>	<u>002</u>	Jan 10, 2019
<u>AB</u>			<u>40MG</u>	<u>A210552</u>	<u>003</u>	Jan 10, 2019
<u>AB</u>		SCIEGEN PHARMS INC	<u>5MG</u>	<u>A208130</u>	<u>001</u>	Jun 29, 2018
<u>AB</u>			<u>20MG</u>	<u>A208130</u>	<u>002</u>	Jun 29, 2018
<u>AB</u>			<u>40MG</u>	<u>A208130</u>	<u>003</u>	Jun 29, 2018
<u>AB</u>		SUNSHINE	<u>5MG</u>	<u>A211049</u>	<u>001</u>	Feb 22, 2019
<u>AB</u>			<u>20MG</u>	<u>A211049</u>	<u>002</u>	Feb 22, 2019
<u>AB</u>			<u>40MG</u>	<u>A211049</u>	<u>003</u>	Feb 22, 2019
<u>AB</u>		UMEDICA	<u>5MG</u>	<u>A207135</u>	<u>001</u>	Jul 18, 2019
<u>AB</u>			<u>20MG</u>	<u>A207135</u>	<u>002</u>	Jul 18, 2019
<u>AB</u>			<u>40MG</u>	<u>A207135</u>	<u>003</u>	Jul 18, 2019
<u>AB</u>		ZYDUS PHARMS	<u>5MG</u>	<u>A205192</u>	<u>001</u>	Apr 24, 2017
<u>AB</u>			<u>20MG</u>	<u>A205192</u>	<u>002</u>	Apr 24, 2017
<u>AB</u>			<u>40MG</u>	<u>A205192</u>	<u>003</u>	Apr 24, 2017

OLODATEROL HYDROCHLORIDE

SPRAY, METERED; INHALATION

STRIVERDI RESPIMAT

+	!	BOEHRINGER INGELHEIM	EQ 0.0025MG BASE/INH	N203108	001	Jul 31, 2014
---	---	-------------------------	----------------------	---------	-----	--------------

PRESCRIPTION DRUG PRODUCT LIST

OLODATEROL HYDROCHLORIDE; TIOTROPIUM BROMIDE

SPRAY, METERED; INHALATION

STIOLTO RESPIMAT

+! BOEHRINGER EQ 0.0025MG BASE/INH; EQ 0.0025MG
INGELHEIM BASE/INH

N206756 001 May 21, 2015

OLOPATADINE HYDROCHLORIDE

SPRAY, METERED; NASAL

OLOPATADINE HYDROCHLORIDE**AB** ! AMNEAL **0.665MG/SPRAY****A210901 001** Jan 28, 2020**AB** APOTEX INC **0.665MG/SPRAY****A091572 001** Oct 08, 2014**AB** PADAGIS ISRAEL **0.665MG/SPRAY****A202853 001** Jan 31, 2017OLSALAZINE SODIUM

CAPSULE; ORAL

DIPENTUM

+! MYLAN SPCLT VIATRIS 250MG

N019715 001 Jul 31, 1990

OLUTASIDENIB

CAPSULE; ORAL

REZLIDHIA

+! RIGEL PHARMS 150MG

N215814 001 Dec 01, 2022

OMADACYCLINE TOSYLATE

POWDER; INTRAVENOUS

NUZYRA

+! PARATEK PHARMS INC EQ 100MG BASE/VIAL

N209817 001 Oct 02, 2018

TABLET; ORAL

NUZYRA

+! PARATEK PHARMS INC EQ 150MG BASE

N209816 001 Oct 02, 2018

OMAVELOXOLONE

CAPSULE; ORAL

SKYCLARYS

+! REATA PHARMS 50MG

N216718 001 Feb 28, 2023

OMEGA-3-ACID ETHYL ESTERS

CAPSULE; ORAL

LOVAZA**AB** +! WOODWARD **1GM CONTAINS AT LEAST 900MG OF THE
ETHYL ESTERS OF OMEGA-3 FATTY ACIDS****N021654 001** Nov 10, 2004OMEGA-3-ACID ETHYL ESTERS**AB** AMNEAL PHARMS **1GM CONTAINS AT LEAST 900MG OF THE
ETHYL ESTERS OF OMEGA-3 FATTY ACIDS****A204940 001** Nov 27, 2015**AB** APOTEX **1GM CONTAINS AT LEAST 900MG OF THE
ETHYL ESTERS OF OMEGA-3 FATTY ACIDS****A090973 001** Sep 30, 2014**AB** ASCENT PHARMS INC **1GM CONTAINS AT LEAST 900MG OF THE
ETHYL ESTERS OF OMEGA-3 FATTY ACIDS****A207420 001** Feb 25, 2019**AB** CHARTWELL **1GM CONTAINS AT LEAST 900MG OF THE
ETHYL ESTERS OF OMEGA-3 FATTY ACIDS****A206455 001** Aug 07, 2019**AB** CSPEC-NBP PHARM **1GM CONTAINS AT LEAST 900MG OF THE
ETHYL ESTERS OF OMEGA-3 FATTY ACIDS****A211979 001** May 12, 2020**AB** GLW **1GM CONTAINS AT LEAST 900MG OF THE
ETHYL ESTERS OF OMEGA-3 FATTY ACIDS****A091028 001** Apr 07, 2014**AB** **1GM CONTAINS AT LEAST 900MG OF THE
ETHYL ESTERS OF OMEGA-3 FATTY ACIDS****A212504 001** Aug 19, 2020**AB** MANKIND PHARMA **1GM CONTAINS AT LEAST 900MG OF THE
ETHYL ESTERS OF OMEGA-3 FATTY ACIDS****A215458 001** Nov 15, 2021**AB** PURACAP PHARM LLC **1GM CONTAINS AT LEAST 900MG OF THE
ETHYL ESTERS OF OMEGA-3 FATTY ACIDS****A210093 001** Jun 15, 2020**AB** SOFGEN PHARMS **1GM CONTAINS AT LEAST 900MG OF THE
ETHYL ESTERS OF OMEGA-3 FATTY ACIDS****A211355 001** Jul 10, 2019**AB** STRIDES SOFTGELS **1GM CONTAINS AT LEAST 900MG OF THE
ETHYL ESTERS OF OMEGA-3 FATTY ACIDS****A203893 001** Sep 19, 2017**AB** SUN PHARM **1GM CONTAINS AT LEAST 900MG OF THE
ETHYL ESTERS OF OMEGA-3 FATTY ACIDS****A210834 001** Jan 09, 2020**AB** WILSHIRE PHARMS INC **1GM CONTAINS AT LEAST 900MG OF THE
ETHYL ESTERS OF OMEGA-3 FATTY ACIDS****A211345 001** Dec 07, 2023OMEPRAZOLE

CAPSULE, DELAYED REL PELLETS; ORAL

OMEPRAZOLE**AB** ACTAVIS LABS FL INC **10MG****A075347 001** May 30, 2008**AB** **20MG****A075347 002** May 30, 2008**AB** **40MG****A075347 003** May 30, 2008**AB** APOTEX **10MG****A076048 001** Oct 22, 2007**AB** **20MG****A076048 002** Oct 22, 2007**AB** **40MG****A076048 003** Jan 21, 2009**AB** AUROBINDO PHARMA **10MG****A203270 001** Aug 19, 2015

PRESCRIPTION DRUG PRODUCT LIST

OMEPRAZOLE

CAPSULE, DELAYED REL PELLETS;ORAL

OMEPRAZOLE

<u>AB</u>		<u>20MG</u>	<u>A203270 002</u>	Aug 19, 2015
<u>AB</u>		<u>40MG</u>	<u>A203270 003</u>	Aug 19, 2015
<u>AB</u>	BRECKENRIDGE	<u>10MG</u>	<u>A203481 001</u>	Jul 03, 2017
<u>AB</u>		<u>20MG</u>	<u>A203481 002</u>	Jul 03, 2017
<u>AB</u>		<u>40MG</u>	<u>A203481 003</u>	Jul 03, 2017
<u>AB</u>	DR REDDYS LABS LTD	<u>10MG</u>	<u>A075576 003</u>	Oct 22, 2007
<u>AB</u>		<u>10MG</u>	<u>A078490 002</u>	Mar 16, 2009
<u>AB</u>		<u>20MG</u>	<u>A075576 002</u>	Oct 22, 2007
<u>AB</u>		<u>20MG</u>	<u>A078490 003</u>	Mar 16, 2009
<u>AB</u>		<u>40MG</u>	<u>A075576 001</u>	Jan 21, 2009
<u>AB</u>		<u>40MG</u>	<u>A078490 001</u>	Apr 17, 2009
<u>AB</u>	GLENMARK PHARMS LTD	<u>10MG</u>	<u>A091672 001</u>	Oct 31, 2014
<u>AB</u>		<u>20MG</u>	<u>A091672 002</u>	Oct 31, 2014
<u>AB</u>		<u>40MG</u>	<u>A091672 003</u>	Oct 31, 2014
<u>AB</u>	HETERO LABS LTD III	<u>10MG</u>	<u>A204012 001</u>	Sep 26, 2019
<u>AB</u>		<u>20MG</u>	<u>A204012 002</u>	Sep 26, 2019
<u>AB</u>	IMPAX LABS	<u>10MG</u>	<u>A075785 001</u>	Oct 22, 2007
<u>AB</u>		<u>20MG</u>	<u>A075785 002</u>	Oct 22, 2007
<u>AB</u>		<u>40MG</u>	<u>A075785 003</u>	Jan 21, 2009
<u>AB</u>	LANNETT CO INC	<u>10MG</u>	<u>A075410 001</u>	Nov 01, 2002
<u>AB</u>		<u>20MG</u>	<u>A075410 002</u>	Nov 01, 2002
<u>AB</u>		<u>40MG</u>	<u>A075410 003</u>	Jan 23, 2009
<u>AB</u>	SANDOZ	<u>10MG</u>	<u>A075757 001</u>	Jan 28, 2003
<u>AB</u>	!	<u>20MG</u>	<u>A075757 002</u>	Jan 28, 2003
<u>AB</u>	!	<u>40MG</u>	<u>A076515 001</u>	Jan 21, 2009
<u>AB</u>	TEVA PHARMS USA	<u>20MG</u>	<u>A204661 001</u>	Jun 13, 2017
<u>AB</u>	XIROMED	<u>10MG</u>	<u>A212977 001</u>	Dec 10, 2020
<u>AB</u>		<u>20MG</u>	<u>A212977 002</u>	Dec 10, 2020
<u>AB</u>		<u>40MG</u>	<u>A212977 003</u>	Dec 27, 2024
<u>AB</u>	ZYDUS PHARMS USA INC	<u>10MG</u>	<u>A091352 001</u>	Nov 19, 2012
<u>AB</u>		<u>20MG</u>	<u>A091352 002</u>	Nov 19, 2012
<u>AB</u>		<u>40MG</u>	<u>A091352 003</u>	Nov 19, 2012

OMEPRAZOLE MAGNESIUM

FOR SUSPENSION, DELAYED RELEASE;ORAL

PRILOSEC

+	COVIS	EQ 2.5MG BASE/PACKET	N022056 001	Mar 20, 2008
+	!	EQ 10MG BASE/PACKET	N022056 002	Mar 20, 2008

OMEPRAZOLE; SODIUM BICARBONATE

CAPSULE;ORAL

OMEPRAZOLE AND SODIUM BICARBONATE

<u>AB</u>	AJANTA PHARMA LTD	<u>20MG;1.1GM</u>	<u>A204228 001</u>	Jul 15, 2016
<u>AB</u>	!	<u>40MG;1.1GM</u>	<u>A204228 002</u>	Jul 15, 2016
<u>AB</u>	ANDA REPOSITORY	<u>20MG;1.1GM</u>	<u>A212587 001</u>	Apr 30, 2020
<u>AB</u>		<u>40MG;1.1GM</u>	<u>A212587 002</u>	Apr 30, 2020
<u>AB</u>	AUROLIFE PHARMA LLC	<u>20MG;1.1GM</u>	<u>A204922 001</u>	Aug 19, 2016
<u>AB</u>		<u>40MG;1.1GM</u>	<u>A204922 002</u>	Aug 19, 2016
<u>AB</u>	DR REDDYS	<u>20MG;1.1GM</u>	<u>A204068 001</u>	Jul 15, 2016
<u>AB</u>		<u>40MG;1.1GM</u>	<u>A204068 002</u>	Jul 15, 2016
<u>AB</u>	SCIEGEN PHARMS INC	<u>20MG;1.1GM</u>	<u>A207476 001</u>	Dec 06, 2016
<u>AB</u>		<u>40MG;1.1GM</u>	<u>A207476 002</u>	Dec 06, 2016
<u>AB</u>	ZYDUS PHARMS	<u>20MG;1.1GM</u>	<u>A203290 001</u>	May 25, 2018
<u>AB</u>		<u>40MG;1.1GM</u>	<u>A203290 002</u>	May 25, 2018

FOR SUSPENSION;ORAL

OMEPRAZOLE AND SODIUM BICARBONATE

<u>AB</u>	AJANTA PHARMA LTD	<u>20MG/PACKET;1.68GM/PACKET</u>	<u>A205545 001</u>	Jul 27, 2016	
<u>AB</u>	!	<u>40MG/PACKET;1.68GM/PACKET</u>	<u>A205545 002</u>	Jul 27, 2016	
<u>AB</u>	STRIDES PHARMA	<u>20MG/PACKET;1.68GM/PACKET</u>	<u>A079182 001</u>	Apr 19, 2013	
<u>AB</u>		<u>40MG/PACKET;1.68GM/PACKET</u>	<u>A079182 002</u>	Apr 19, 2013	
	KONVOMEPEP				
+	!	AZURITY	2MG/ML; 84MG/ML	N213593 001	Aug 30, 2022

ONDANSETRON

TABLET, ORALLY DISINTEGRATING;ORAL

ONDANSETRON

<u>AB</u>	AUROBINDO PHARMA	<u>4MG</u>	<u>A090469 001</u>	Apr 12, 2010
<u>AB</u>	!	<u>8MG</u>	<u>A090469 002</u>	Apr 12, 2010
<u>AB</u>	CHARTWELL MOLECULES	<u>4MG</u>	<u>A077406 003</u>	Dec 26, 2006
<u>AB</u>		<u>8MG</u>	<u>A077406 004</u>	Dec 26, 2006

PRESCRIPTION DRUG PRODUCT LIST

ONDANSETRON

TABLET, ORALLY DISINTEGRATING;ORAL

ONDANSETRON

<u>AB</u>	GLENMARK PHARMS LTD	<u>4MG</u>	<u>A078152</u>	<u>001</u>	Jun 27, 2007
<u>AB</u>		<u>8MG</u>	<u>A078152</u>	<u>002</u>	Jun 27, 2007
<u>AB</u>	SANDOZ	<u>4MG</u>	<u>A078050</u>	<u>001</u>	Aug 13, 2007
<u>AB</u>		<u>8MG</u>	<u>A078050</u>	<u>002</u>	Aug 13, 2007
<u>AB</u>	SUN PHARM INDS	<u>4MG</u>	<u>A077557</u>	<u>001</u>	Aug 02, 2007
<u>AB</u>		<u>8MG</u>	<u>A077557</u>	<u>002</u>	Aug 02, 2007
!	CHARTWELL MOLECULES	16MG	A077406	001	Dec 26, 2006
!		24MG	A077406	002	Dec 26, 2006

ONDANSETRON HYDROCHLORIDE

INJECTABLE;INJECTION

ONDANSETRON HYDROCHLORIDE

<u>AP</u>	ACCORD HLTHCARE	<u>EQ 2MG BASE/ML</u>	<u>A206846</u>	<u>001</u>	Jul 13, 2015
<u>AP</u>	FRESENIUS KABI USA	<u>EQ 2MG BASE/ML</u>	<u>A076974</u>	<u>001</u>	Dec 26, 2006
<u>AP</u>	GLAND PHARMA LTD	<u>EQ 2MG BASE/ML</u>	<u>A079224</u>	<u>001</u>	Sep 25, 2009
<u>AP</u>	!	<u>EQ 2MG BASE/ML</u>	<u>A090648</u>	<u>001</u>	Jun 15, 2012
<u>AP</u>	HIKMA	<u>EQ 2MG BASE/ML</u>	<u>A076967</u>	<u>001</u>	Dec 26, 2006
<u>AP</u>		<u>EQ 2MG BASE/ML</u>	<u>A077365</u>	<u>001</u>	Dec 26, 2006
<u>AP</u>	HIKMA FARMACEUTICA	<u>EQ 2MG BASE/ML</u>	<u>A076781</u>	<u>001</u>	Dec 26, 2006
<u>AP</u>	HOSPIRA	<u>EQ 2MG BASE/ML</u>	<u>A077473</u>	<u>001</u>	Dec 26, 2006
<u>AP</u>	!	<u>EQ 2MG BASE/ML</u>	<u>A203711</u>	<u>001</u>	Sep 08, 2014
<u>AP</u>	SANDOZ	<u>EQ 2MG BASE/ML</u>	<u>A077430</u>	<u>001</u>	Jun 27, 2007
<u>AP</u>	WOCKHARDT	<u>EQ 2MG BASE/ML</u>	<u>A077577</u>	<u>001</u>	Dec 26, 2006

ONDANSETRON HYDROCHLORIDE PRESERVATIVE FREE

<u>AP</u>	!	BAXTER HLTHCARE	<u>EQ 2MG BASE/ML</u>	<u>A078287</u>	<u>001</u>	Feb 22, 2013
		CORP				
<u>AP</u>		FRESENIUS KABI USA	<u>EQ 2MG BASE/ML</u>	<u>A076972</u>	<u>001</u>	Dec 26, 2006
<u>AP</u>			<u>EQ 2MG BASE/ML</u>	<u>A202253</u>	<u>001</u>	Jul 19, 2013
<u>AP</u>		HIKMA	<u>EQ 2MG BASE/ML</u>	<u>A077011</u>	<u>001</u>	Dec 26, 2006
<u>AP</u>			<u>EQ 2MG BASE/ML</u>	<u>A077541</u>	<u>001</u>	Dec 26, 2006
<u>AP</u>		HOSPIRA	<u>EQ 2MG BASE/ML</u>	<u>A077548</u>	<u>001</u>	Dec 26, 2006
<u>AP</u>		SANDOZ	<u>EQ 2MG BASE/ML</u>	<u>A077551</u>	<u>001</u>	Jun 27, 2007
<u>AP</u>		WOCKHARDT	<u>EQ 2MG BASE/ML</u>	<u>A077716</u>	<u>001</u>	Dec 26, 2006

SOLUTION;ORAL

ONDANSETRON HYDROCHLORIDE

<u>AA</u>	AMNEAL PHARMS	<u>EQ 4MG BASE/5ML</u>	<u>A091483</u>	<u>001</u>	Jan 31, 2011	
<u>AA</u>	AUROBINDO PHARMA	<u>EQ 4MG BASE/5ML</u>	<u>A078776</u>	<u>001</u>	Nov 28, 2007	
<u>AA</u>	CHARTWELL MOLECULAR	<u>EQ 4MG BASE/5ML</u>	<u>A091342</u>	<u>001</u>	Jan 27, 2011	
<u>AA</u>	!	HIKMA	<u>EQ 4MG BASE/5ML</u>	<u>A076960</u>	<u>001</u>	Dec 26, 2006

TABLET;ORAL

ONDANSETRON HYDROCHLORIDE

<u>AB</u>	APOTEX	<u>EQ 4MG BASE</u>	<u>A077306</u>	<u>001</u>	Jun 25, 2007
<u>AB</u>		<u>EQ 8MG BASE</u>	<u>A077306</u>	<u>002</u>	Jun 25, 2007
<u>AB</u>	AUROBINDO PHARMA	<u>EQ 4MG BASE</u>	<u>A078539</u>	<u>001</u>	Jul 31, 2007
<u>AB</u>		<u>EQ 8MG BASE</u>	<u>A078539</u>	<u>002</u>	Jul 31, 2007
<u>AB</u>	CHARTWELL MOLECULES	<u>EQ 4MG BASE</u>	<u>A077303</u>	<u>001</u>	Jun 25, 2007
<u>AB</u>		<u>EQ 8MG BASE</u>	<u>A077303</u>	<u>002</u>	Jun 25, 2007
<u>AB</u>	DR REDDYS LABS LTD	<u>EQ 4MG BASE</u>	<u>A076183</u>	<u>003</u>	Dec 26, 2006
<u>AB</u>		<u>EQ 8MG BASE</u>	<u>A076183</u>	<u>002</u>	Dec 26, 2006
<u>AB</u>	GLENMARK PHARMS LTD	<u>EQ 4MG BASE</u>	<u>A077535</u>	<u>001</u>	Jun 25, 2007
<u>AB</u>		<u>EQ 8MG BASE</u>	<u>A077535</u>	<u>002</u>	Jun 25, 2007
<u>AB</u>	NATCO PHARMA LTD	<u>EQ 4MG BASE</u>	<u>A077851</u>	<u>001</u>	Jun 25, 2007
<u>AB</u>		<u>EQ 8MG BASE</u>	<u>A077851</u>	<u>002</u>	Jun 25, 2007
!	CHARTWELL MOLECULES	EQ 24MG BASE	A077303	004	Jun 25, 2007

OPICAPONE

CAPSULE;ORAL

ONGENTYS

+	AMNEAL	25MG	N212489	001	Apr 24, 2020
+	!	50MG	N212489	002	Apr 24, 2020

ORITAVANCIN DIPHOSPHATE

POWDER;INTRAVENOUS

KIMYRSA

+	!	MELINTA THERAP	EQ 1.2GM BASE/VIAL	N214155	001	Mar 12, 2021
		ORBACTIV				
+	!	MELINTA THERAP	EQ 400MG BASE/VIAL	N206334	001	Aug 06, 2014

PRESCRIPTION DRUG PRODUCT LIST

ORLISTAT

CAPSULE; ORAL

XENICAL

+! CHEPLAPHARM

120MG

N020766 001 Apr 23, 1999

ORPHENADRINE CITRATE

INJECTABLE; INJECTION

ORPHENADRINE CITRATEAP HIKMA30MG/MLA040463 001 Mar 04, 2003AP ! RISING30MG/MLA040484 001 May 24, 2006AP SAGENT PHARMS30MG/MLA090585 001 Aug 30, 2011AP WATSON LABS30MG/MLA084779 001 Mar 15, 1982

TABLET, EXTENDED RELEASE; ORAL

ORPHENADRINE CITRATEAB ! LUPIN100MGA040284 001 Jun 19, 1998AB RISING100MGA040249 001 Jan 29, 1999AB SANDOZ100MGA040327 001 Feb 15, 2000AB UNICHEM100MGA091158 001 Jul 27, 2012OSELTAMIVIR PHOSPHATE

CAPSULE; ORAL

OSELTAMIVIR PHOSPHATEAB ALEMBICEQ 30MG BASEA211823 001 Jun 24, 2019ABEQ 45MG BASEA211823 002 Jun 24, 2019ABEQ 75MG BASEA211823 003 Jun 24, 2019AB AMNEAL PHARMSEQ 30MG BASEA209093 001 May 17, 2017ABEQ 45MG BASEA209093 002 May 17, 2017ABEQ 75MG BASEA209093 003 May 17, 2017AB CEDIPROF INCEQ 30MG BASEA217451 001 Dec 03, 2024ABEQ 45MG BASEA217451 002 Dec 03, 2024ABEQ 75MG BASEA217451 003 Dec 03, 2024AB EPIC PHARMA LLCEQ 30MG BASEA215208 001 Oct 01, 2021ABEQ 45MG BASEA215208 002 Oct 01, 2021ABEQ 75MG BASEA215208 003 Oct 01, 2021AB HAINAN POLYEQ 75MG BASEA218009 001 Nov 24, 2023AB HETERO LABS LTD IIIEQ 30MG BASEA209438 001 Feb 23, 2018ABEQ 45MG BASEA209438 002 Feb 23, 2018ABEQ 75MG BASEA209438 003 Feb 23, 2018AB INVAGEN PHARMSEQ 30MG BASEA217467 001 Oct 03, 2023ABEQ 45MG BASEA217467 002 Oct 03, 2023ABEQ 75MG BASEA217467 003 Oct 03, 2023AB LAURUSEQ 30MG BASEA218565 001 May 01, 2024ABEQ 45MG BASEA218565 002 May 01, 2024ABEQ 75MG BASEA218565 003 May 01, 2024AB LUPINEQ 30MG BASEA208348 001 Jan 09, 2018ABEQ 45MG BASEA208348 002 Jan 09, 2018ABEQ 75MG BASEA208348 003 Jan 09, 2018AB MACLEODS PHARMS LTDEQ 30MG BASEA207211 001 Sep 14, 2017ABEQ 45MG BASEA207211 002 Sep 14, 2017ABEQ 75MG BASEA207211 003 Sep 14, 2017AB MSNEQ 30MG BASEA212544 001 May 20, 2020ABEQ 45MG BASEA212544 002 May 20, 2020ABEQ 75MG BASEA212544 003 May 20, 2020AB NANJING DAOQUNEQ 30MG BASEA217133 001 Oct 30, 2024ABEQ 75MG BASEA217133 002 Oct 30, 2024AB NATCOEQ 30MG BASEA202595 001 Aug 03, 2016ABEQ 45MG BASEA202595 002 Aug 03, 2016ABEQ 75MG BASEA202595 003 Aug 03, 2016AB STRIDES PHARMAEQ 30MG BASEA209421 001 Jun 08, 2018ABEQ 45MG BASEA209421 002 Jun 08, 2018ABEQ 75MG BASEA209421 003 Jun 08, 2018AB SUNSHINEEQ 30MG BASEA212739 001 Mar 04, 2020ABEQ 45MG BASEA212739 002 Mar 04, 2020ABEQ 75MG BASEA212739 003 Mar 04, 2020AB ZYDUS PHARMSEQ 30MG BASEA208578 001 Feb 24, 2017ABEQ 45MG BASEA208578 002 Feb 24, 2017ABEQ 75MG BASEA208578 003 Feb 24, 2017TAMIFLUAB + ROCHEEQ 30MG BASEN021087 003 Jul 02, 2007AB +EQ 45MG BASEN021087 002 Jul 02, 2007AB +!EQ 75MG BASEN021087 001 Oct 27, 1999

PRESCRIPTION DRUG PRODUCT LIST

OSELTAMIVIR PHOSPHATE

FOR SUSPENSION; ORAL

OSELTAMIVIR PHOSPHATE

AB	AJANTA PHARMA LTD	EQ 6MG BASE/ML	A212784 001	May 27, 2020
AB	ALVOGEN	EQ 6MG BASE/ML	A208823 001	Oct 31, 2017
AB	AMNEAL PHARMS NY	EQ 6MG BASE/ML	A210186 001	Feb 27, 2018
AB	AUROBINDO PHARMA LTD	EQ 6MG BASE/ML	A218131 001	May 10, 2024
AB	EPIC PHARMA LLC	EQ 6MG BASE/ML	A215538 001	Feb 14, 2023
AB	HETERO LABS LTD V	EQ 6MG BASE/ML	A209590 001	Mar 28, 2022
AB	INVAGEN PHARMS	EQ 6MG BASE/ML	A212858 001	Aug 30, 2021
AB	LUPIN	EQ 6MG BASE/ML	A208347 001	Feb 20, 2018
AB	MSN	EQ 6MG BASE/ML	A215313 001	May 02, 2022
AB	ORYZA	EQ 6MG BASE/ML	A217303 001	Jul 02, 2024
AB	STRIDES PHARMA	EQ 6MG BASE/ML	A211894 001	Jan 13, 2022
AB	SUNSHINE	EQ 6MG BASE/ML	A213594 001	Jan 05, 2022
AB	TEVA PHARMS USA	EQ 6MG BASE/ML	A211125 001	Feb 27, 2019
AB	ZYDUS PHARMS	EQ 6MG BASE/ML	A209113 001	Sep 14, 2017

TAMIFLU

AB	+ ! ROCHE	EQ 6MG BASE/ML	N021246 002	Mar 21, 2011
-----------	------------------	-----------------------	--------------------	--------------

OSILODROSTAT PHOSPHATE

TABLET; ORAL

ISTURISA

+	RECORDATI RARE	EQ 1MG BASE	N212801 001	Mar 06, 2020
+		EQ 5MG BASE	N212801 002	Mar 06, 2020

OSIMERTINIB MESYLATE

TABLET; ORAL

TAGRISSO

+	ASTRAZENECA	EQ 40MG BASE	N208065 001	Nov 13, 2015
+ !		EQ 80MG BASE	N208065 002	Nov 13, 2015

OSPEMIFENE

TABLET; ORAL

OSPEMIFENE

AB	HETERO LABS LTD V	60MG	A215574 001	Feb 13, 2024
AB	+ ! DUCHESNAY	60MG	N203505 001	Feb 26, 2013

OTESECONAZOLE

CAPSULE; ORAL

VIVJOA

+ !	MYCOVIA PHARMS	150MG	N215888 001	Apr 26, 2022
------------	----------------	-------	-------------	--------------

OXACILLIN SODIUM

INJECTABLE; INJECTION

OXACILLIN SODIUM

AP	EUGIA PHARMA SPECLTS	EQ 1GM BASE/VIAL	A201539 001	Jan 18, 2013
AP		EQ 2GM BASE/VIAL	A201539 002	Jan 18, 2013
AP		EQ 10GM BASE/VIAL	A201538 001	Jan 18, 2013
AP	! FRESENIUS KABI USA	EQ 1GM BASE/VIAL	A206198 001	Jul 20, 2020
AP	!	EQ 2GM BASE/VIAL	A206198 002	Jul 20, 2020
AP		EQ 10GM BASE/VIAL	A206199 001	Jul 27, 2020
AP	! SAGENT PHARMS	EQ 10GM BASE/VIAL	A091245 001	Mar 30, 2012
AP	STERISCIENCE SPECLTS	EQ 2GM BASE/VIAL	A091486 002	Aug 25, 2014
AP	WOCKHARDT BIO AG	EQ 1GM BASE/VIAL	A207147 001	Jul 31, 2017
AP		EQ 2GM BASE/VIAL	A207147 002	Jul 31, 2017
AP		EQ 10GM BASE/VIAL	A207148 001	Nov 24, 2017
	BACTOCILL IN PLASTIC CONTAINER			
+ !	BAXTER HLTHCARE	EQ 20MG BASE/ML	N050640 001	Oct 26, 1989
+ !		EQ 40MG BASE/ML	N050640 002	Oct 26, 1989

OXALIPLATIN

INJECTABLE; INTRAVENOUS

OXALIPLATIN

AP	ACCORD HLTHCARE	50MG/10ML (5MG/ML)	A207474 001	Mar 21, 2017
AP		100MG/20ML (5MG/ML)	A207474 002	Mar 21, 2017
AP	ACTAVIS	50MG/10ML (5MG/ML)	A204880 001	Mar 05, 2018
AP		100MG/20ML (5MG/ML)	A204880 002	Mar 05, 2018
AP	FRESENIUS KABI USA	50MG/10ML (5MG/ML)	A078811 001	Jun 10, 2010
AP		100MG/20ML (5MG/ML)	A078811 002	Jun 10, 2010
AP		50MG/VIAL	A078819 001	Jun 02, 2010
AP		50MG/10ML (5MG/ML)	A090030 001	Jan 31, 2017

PRESCRIPTION DRUG PRODUCT LIST

OXALIPLATIN

INJECTABLE; INTRAVENOUS

OXALIPLATIN

<u>AP</u>		<u>100MG/VIAL</u>	<u>A078819 002</u>	Jun 02, 2010	
<u>AP</u>		<u>100MG/20ML (5MG/ML)</u>	<u>A090030 002</u>	Jan 31, 2017	
<u>AP</u>	!	GLAND	<u>50MG/10ML (5MG/ML)</u>	<u>A207325 001</u>	Feb 10, 2017
<u>AP</u>	!		<u>100MG/20ML (5MG/ML)</u>	<u>A207325 002</u>	Feb 10, 2017
<u>AP</u>		GLAND PHARMA LTD	<u>50MG/VIAL</u>	<u>A207385 001</u>	May 23, 2017
<u>AP</u>			<u>100MG/VIAL</u>	<u>A207385 002</u>	May 23, 2017
<u>AP</u>		HENGRUI PHARMA	<u>50MG/10ML (5MG/ML)</u>	<u>A203869 001</u>	Jun 18, 2014
<u>AP</u>			<u>100MG/20ML (5MG/ML)</u>	<u>A203869 002</u>	Jun 18, 2014
<u>AP</u>		HOSPIRA WORLDWIDE	<u>50MG/10ML (5MG/ML)</u>	<u>A078813 001</u>	Aug 07, 2009
<u>AP</u>			<u>100MG/20ML (5MG/ML)</u>	<u>A078813 002</u>	Aug 07, 2009
<u>AP</u>		MEITHEAL	<u>50MG/10ML (5MG/ML)</u>	<u>A217348 001</u>	Jan 11, 2024
<u>AP</u>			<u>100MG/20ML (5MG/ML)</u>	<u>A217348 002</u>	Jan 11, 2024
<u>AP</u>		MYLAN LABS LTD	<u>50MG/10ML (5MG/ML)</u>	<u>A091358 001</u>	Aug 07, 2012
<u>AP</u>			<u>100MG/20ML (5MG/ML)</u>	<u>A091358 002</u>	Aug 07, 2012
<u>AP</u>		NOVAST LABS	<u>50MG/10ML (5MG/ML)</u>	<u>A207562 001</u>	Oct 16, 2018
<u>AP</u>			<u>100MG/20ML (5MG/ML)</u>	<u>A207562 002</u>	Oct 16, 2018
<u>AP</u>		QILU PHARM HAINAN	<u>50MG/10ML (5MG/ML)</u>	<u>A204368 001</u>	Jun 07, 2016
<u>AP</u>			<u>50MG/VIAL</u>	<u>A204616 001</u>	May 11, 2016
<u>AP</u>			<u>100MG/20ML (5MG/ML)</u>	<u>A204368 002</u>	Jun 07, 2016
<u>AP</u>			<u>100MG/VIAL</u>	<u>A204616 002</u>	May 11, 2016
<u>AP</u>		SANDOZ	<u>50MG/10ML (5MG/ML)</u>	<u>A078817 001</u>	Jan 24, 2011
<u>AP</u>			<u>100MG/20ML (5MG/ML)</u>	<u>A078817 002</u>	Jan 24, 2011
<u>AP</u>	+	TEVA PHARMS	<u>50MG/10ML (5MG/ML)</u>	<u>N022160 001</u>	Aug 07, 2009
<u>AP</u>	+		<u>100MG/20ML (5MG/ML)</u>	<u>N022160 002</u>	Aug 07, 2009
<u>AP</u>	!	QILU PHARM HAINAN	<u>200MG/40ML (5MG/ML)</u>	<u>A204368 003</u>	Jun 07, 2016

OXAPROZIN

CAPSULE; ORAL

COXANTO

+! SOLUBIOMIX

300MG

N217927 001 Oct 20, 2023

TABLET; ORAL

DAYPRO

<u>AB</u>	+	PFIZER	<u>600MG</u>	<u>N018841 004</u>	Oct 29, 1992
-----------	---	--------	--------------	--------------------	--------------

OXAPROZIN

<u>AB</u>		AMNEAL PHARMS CO	<u>600MG</u>	<u>A208633 001</u>	May 04, 2017
<u>AB</u>		CHARTWELL	<u>600MG</u>	<u>A075987 001</u>	Sep 02, 2004
<u>AB</u>		DR REDDYS LABS LTD	<u>600MG</u>	<u>A075855 001</u>	Jan 31, 2001
<u>AB</u>		PANGEA	<u>600MG</u>	<u>A075845 001</u>	Jan 31, 2001

OXAZEPAM

CAPSULE; ORAL

OXAZEPAM

<u>AB</u>		ACTAVIS ELIZABETH	<u>10MG</u>	<u>A072253 002</u>	Apr 14, 1988
<u>AB</u>			<u>15MG</u>	<u>A072253 003</u>	Apr 14, 1988
<u>AB</u>	!		<u>30MG</u>	<u>A072253 001</u>	Apr 14, 1988
<u>AB</u>		EPIC PHARMA LLC	<u>10MG</u>	<u>A071813 001</u>	Apr 19, 1988
<u>AB</u>			<u>15MG</u>	<u>A071756 001</u>	Apr 19, 1988
<u>AB</u>			<u>30MG</u>	<u>A071814 001</u>	Apr 19, 1988
<u>AB</u>		TRUPHARMA	<u>10MG</u>	<u>A071026 002</u>	Aug 10, 1987
<u>AB</u>			<u>15MG</u>	<u>A071026 003</u>	Aug 10, 1987
<u>AB</u>			<u>30MG</u>	<u>A071026 001</u>	Aug 10, 1987

OXCARBAZEPINE

SUSPENSION; ORAL

OXCARBAZEPINE

<u>AB</u>		ALKEM LABS LTD	<u>300MG/5ML</u>	<u>A213183 001</u>	Mar 18, 2023
<u>AB</u>		AMNEAL PHARMS	<u>300MG/5ML</u>	<u>A202961 001</u>	Sep 17, 2012
<u>AB</u>		AUCTA	<u>300MG/5ML</u>	<u>A215332 001</u>	Nov 30, 2022
<u>AB</u>		BIOCON PHARMA	<u>300MG/5ML</u>	<u>A218278 001</u>	Mar 01, 2024
<u>AB</u>		BIONPHARMA	<u>300MG/5ML</u>	<u>A209652 001</u>	Nov 04, 2022
<u>AB</u>		CHARTWELL RX	<u>300MG/5ML</u>	<u>A212428 001</u>	Jun 21, 2021
<u>AB</u>		HETERO LABS LTD III	<u>300MG/5ML</u>	<u>A216749 001</u>	Oct 20, 2023
<u>AB</u>		KANCHAN HLTHCARE	<u>300MG/5ML</u>	<u>A217782 001</u>	Feb 14, 2024
<u>AB</u>		RUBICON	<u>300MG/5ML</u>	<u>A215726 001</u>	Aug 30, 2022
<u>AB</u>		SUN PHARM INDS LTD	<u>300MG/5ML</u>	<u>A078734 001</u>	Jun 26, 2009

TRILEPTAL

<u>AB</u>	+	NOVARTIS	<u>300MG/5ML</u>	<u>N021285 001</u>	May 25, 2001
-----------	---	----------	------------------	--------------------	--------------

TABLET; ORAL

OXCARBAZEPINE

<u>AB</u>		ANI PHARMS	<u>150MG</u>	<u>A078005 001</u>	Dec 11, 2007
<u>AB</u>			<u>300MG</u>	<u>A078005 002</u>	Dec 11, 2007

PRESCRIPTION DRUG PRODUCT LIST

OXCARBAZEPINE

TABLET; ORAL

OXCARBAZEPINE

<u>AB</u>		<u>600MG</u>	<u>A078005 003</u>	Dec 11, 2007
<u>AB</u>	ANNORA PHARMA	<u>150MG</u>	<u>A215939 001</u>	Jan 11, 2022
<u>AB</u>		<u>300MG</u>	<u>A215939 002</u>	Jan 11, 2022
<u>AB</u>		<u>600MG</u>	<u>A215939 003</u>	Jan 11, 2022
<u>AB</u>	BRECKENRIDGE PHARM	<u>150MG</u>	<u>A078069 001</u>	Jan 11, 2008
<u>AB</u>		<u>300MG</u>	<u>A078069 002</u>	Jan 11, 2008
<u>AB</u>		<u>600MG</u>	<u>A078069 003</u>	Jan 11, 2008
<u>AB</u>	GLENMARK PHARMS LTD	<u>150MG</u>	<u>A077802 001</u>	Oct 09, 2007
<u>AB</u>		<u>300MG</u>	<u>A077802 002</u>	Oct 09, 2007
<u>AB</u>		<u>600MG</u>	<u>A077802 003</u>	Oct 09, 2007
<u>AB</u>	RUBICON	<u>150MG</u>	<u>A077747 001</u>	Apr 09, 2008
<u>AB</u>		<u>150MG</u>	<u>A207717 001</u>	Mar 05, 2024
<u>AB</u>		<u>300MG</u>	<u>A077747 002</u>	Apr 09, 2008
<u>AB</u>		<u>300MG</u>	<u>A207717 002</u>	Mar 05, 2024
<u>AB</u>		<u>600MG</u>	<u>A077747 003</u>	Apr 09, 2008
<u>AB</u>		<u>600MG</u>	<u>A207717 003</u>	Mar 05, 2024
<u>AB</u>	SUN PHARM INDS	<u>150MG</u>	<u>A077794 001</u>	Oct 09, 2007
<u>AB</u>		<u>300MG</u>	<u>A077794 002</u>	Oct 09, 2007
<u>AB</u>		<u>600MG</u>	<u>A077794 003</u>	Oct 09, 2007
<u>AB</u>	TARO	<u>150MG</u>	<u>A077801 001</u>	Nov 15, 2007
<u>AB</u>		<u>300MG</u>	<u>A077801 002</u>	Nov 15, 2007
<u>AB</u>		<u>600MG</u>	<u>A077801 003</u>	Nov 15, 2007

TRILEPTAL

<u>AB</u>	+ NOVARTIS	<u>150MG</u>	<u>N021014 001</u>	Jan 14, 2000
<u>AB</u>	+	<u>300MG</u>	<u>N021014 002</u>	Jan 14, 2000
<u>AB</u>	+	<u>600MG</u>	<u>N021014 003</u>	Jan 14, 2000

TABLET, EXTENDED RELEASE; ORAL

OXCARBAZEPINE

<u>AB</u>	APOTEX	<u>150MG</u>	<u>A213369 001</u>	Jul 13, 2023
<u>AB</u>		<u>300MG</u>	<u>A213369 002</u>	Jul 13, 2023
<u>AB</u>		<u>600MG</u>	<u>A213369 003</u>	Jul 13, 2023

OXTELLAR XR

<u>AB</u>	+ SUPERNUS PHARMS	<u>150MG</u>	<u>N202810 001</u>	Oct 19, 2012
<u>AB</u>	+	<u>300MG</u>	<u>N202810 002</u>	Oct 19, 2012
<u>AB</u>	+	<u>600MG</u>	<u>N202810 003</u>	Oct 19, 2012

OXICONAZOLE NITRATE

CREAM; TOPICAL

OXICONAZOLE NITRATE

<u>AB</u>	TARO	<u>EQ 1% BASE</u>	<u>A205076 001</u>	Mar 07, 2016
<u>AB</u>	+	<u>EQ 1% BASE</u>	<u>N019828 001</u>	Dec 30, 1988

LOTION; TOPICAL

OXISTAT

+	ANI PHARMS	<u>EQ 1% BASE</u>	<u>N020209 001</u>	Sep 30, 1992
---	------------	-------------------	--------------------	--------------

OXYBUTYNIN

FILM, EXTENDED RELEASE; TRANSDERMAL

OXYTROL

+	ALLERGAN	<u>3.9MG/24HR</u>	<u>N021351 002</u>	Feb 26, 2003
---	----------	-------------------	--------------------	--------------

OXYBUTYNIN CHLORIDE

SYRUP; ORAL

OXYBUTYNIN CHLORIDE

<u>AA</u>	CHARTWELL RX	<u>5MG/5ML</u>	<u>A075039 001</u>	Jan 29, 1999
<u>AA</u>	! LANNETT CO INC	<u>5MG/5ML</u>	<u>A074520 001</u>	Mar 29, 1996

TABLET; ORAL

OXYBUTYNIN CHLORIDE

<u>AB</u>	ABHAI LLC	<u>5MG</u>	<u>A209335 001</u>	Dec 22, 2017
<u>AB</u>	BEXIMCO PHARMS USA	<u>5MG</u>	<u>A213550 001</u>	Jul 14, 2022
<u>AB</u>	LEADING	<u>5MG</u>	<u>A212798 001</u>	Apr 06, 2020
<u>AB</u>	NOVAST LABS	<u>5MG</u>	<u>A210611 001</u>	Oct 30, 2019
<u>AB</u>	NOVITIUM PHARMA	<u>5MG</u>	<u>A209823 001</u>	Oct 23, 2017
<u>AB</u>	RISING	<u>5MG</u>	<u>A209025 001</u>	Dec 21, 2017
<u>AB</u>	! STRIDES PHARMA	<u>5MG</u>	<u>A075079 001</u>	Oct 31, 1997
<u>AB</u>		<u>5MG</u>	<u>A208165 001</u>	Dec 17, 2020
<u>AB</u>	TEVA PHARMS USA	<u>5MG</u>	<u>A071655 001</u>	Nov 14, 1988
<u>AB</u>	TRUPHARMA	<u>5MG</u>	<u>A210125 001</u>	Sep 06, 2018
<u>AB</u>	UPSHER SMITH LABS	<u>5MG</u>	<u>A074625 001</u>	Jul 31, 1996
!	RISING	<u>2.5MG</u>	<u>A209025 002</u>	Feb 07, 2023

PRESCRIPTION DRUG PRODUCT LIST

OXYBUTYNIN CHLORIDE

TABLET, EXTENDED RELEASE;ORAL

OXYBUTYNIN CHLORIDE

<u>AB</u>	ACCORD HLTHCARE	<u>5MG</u>	<u>A207138</u>	<u>001</u>	Feb 29, 2016
<u>AB</u>		<u>10MG</u>	<u>A207138</u>	<u>002</u>	Feb 29, 2016
<u>AB</u>	!	<u>15MG</u>	<u>A207138</u>	<u>003</u>	Feb 29, 2016
<u>AB</u>	AJANTA PHARMA LTD	<u>5MG</u>	<u>A211655</u>	<u>001</u>	Feb 28, 2019
<u>AB</u>		<u>10MG</u>	<u>A211655</u>	<u>002</u>	Feb 28, 2019
<u>AB</u>		<u>15MG</u>	<u>A211655</u>	<u>003</u>	Feb 28, 2019
<u>AB</u>	AMNEAL PHARMS	<u>5MG</u>	<u>A204010</u>	<u>001</u>	Nov 23, 2015
<u>AB</u>		<u>10MG</u>	<u>A204010</u>	<u>002</u>	Nov 23, 2015
<u>AB</u>		<u>15MG</u>	<u>A204010</u>	<u>003</u>	Nov 23, 2015
<u>AB</u>	BIONPHARMA	<u>5MG</u>	<u>A210717</u>	<u>001</u>	Dec 17, 2019
<u>AB</u>		<u>10MG</u>	<u>A210717</u>	<u>002</u>	Dec 17, 2019
<u>AB</u>		<u>15MG</u>	<u>A210717</u>	<u>003</u>	Dec 17, 2019
<u>AB</u>	OSMOTICA PHARM US	<u>5MG</u>	<u>A078503</u>	<u>001</u>	Feb 04, 2009
<u>AB</u>		<u>10MG</u>	<u>A078503</u>	<u>002</u>	Feb 04, 2009
<u>AB</u>		<u>15MG</u>	<u>A078503</u>	<u>003</u>	Feb 04, 2009
<u>AB</u>	RUBICON	<u>5MG</u>	<u>A214415</u>	<u>001</u>	Oct 27, 2020
<u>AB</u>		<u>10MG</u>	<u>A214415</u>	<u>002</u>	Oct 27, 2020
<u>AB</u>		<u>15MG</u>	<u>A214415</u>	<u>003</u>	Oct 27, 2020
<u>AB</u>	UNIQUE	<u>5MG</u>	<u>A206121</u>	<u>001</u>	May 27, 2016
<u>AB</u>		<u>10MG</u>	<u>A206121</u>	<u>002</u>	May 27, 2016
<u>AB</u>		<u>15MG</u>	<u>A206121</u>	<u>003</u>	May 27, 2016
<u>AB</u>	ZYDUS PHARMS	<u>5MG</u>	<u>A202332</u>	<u>001</u>	Jun 26, 2017
<u>AB</u>		<u>10MG</u>	<u>A202332</u>	<u>002</u>	Jun 26, 2017
<u>AB</u>		<u>15MG</u>	<u>A202332</u>	<u>003</u>	Jun 26, 2017

OXYCODONE

CAPSULE, EXTENDED RELEASE;ORAL

XTAMPZA ER

+	COLLEGIUM PHARM INC	9MG	N208090	001	Apr 26, 2016
+		13.5MG	N208090	002	Apr 26, 2016
+		18MG	N208090	003	Apr 26, 2016
+		27MG	N208090	004	Apr 26, 2016
+	!	36MG	N208090	005	Apr 26, 2016

OXYCODONE HYDROCHLORIDE

CAPSULE;ORAL

OXYCODONE HYDROCHLORIDE

<u>AB</u>	ANI PHARMS	<u>5MG</u>	<u>A205177</u>	<u>001</u>	Mar 31, 2016
<u>AB</u>	AVANTHI INC	<u>5MG</u>	<u>A202773</u>	<u>001</u>	Aug 17, 2015
<u>AB</u>	+!	GENUS LIFESCIENCES	<u>N200534</u>	<u>001</u>	Oct 20, 2010
<u>AB</u>	NOVEL LABS INC	<u>5MG</u>	<u>A204752</u>	<u>001</u>	Aug 24, 2015

SOLUTION;ORAL

OXYCODONE HYDROCHLORIDE

<u>AA</u>	ABHAI LLC	<u>5MG/5ML</u>	<u>A208593</u>	<u>001</u>	Jul 21, 2017
<u>AA</u>		<u>100MG/5ML</u>	<u>A208593</u>	<u>002</u>	Jul 21, 2017
<u>AA</u>	ALKEM LABS LTD	<u>5MG/5ML</u>	<u>A211748</u>	<u>001</u>	Feb 07, 2019
<u>AA</u>		<u>100MG/5ML</u>	<u>A211749</u>	<u>001</u>	Feb 04, 2019
<u>AA</u>	ANI PHARMS	<u>5MG/5ML</u>	<u>A204979</u>	<u>001</u>	Jun 01, 2015
<u>AA</u>	ASCENT PHARMS INC	<u>5MG/5ML</u>	<u>A209021</u>	<u>001</u>	Nov 09, 2017
<u>AA</u>		<u>100MG/5ML</u>	<u>A209021</u>	<u>002</u>	Nov 09, 2017
<u>AA</u>	+	GENUS LIFESCIENCES	<u>N200535</u>	<u>002</u>	Aug 22, 2013
<u>AA</u>	+	!	<u>N200535</u>	<u>001</u>	Oct 20, 2010
<u>AA</u>	!	HIKMA	<u>A204037</u>	<u>001</u>	Jul 15, 2013
<u>AA</u>	PHARM ASSOC	<u>5MG/5ML</u>	<u>A206914</u>	<u>001</u>	Feb 01, 2019
<u>AA</u>		<u>100MG/5ML</u>	<u>A206822</u>	<u>001</u>	Aug 15, 2017
<u>AA</u>	QUAGEN	<u>5MG/5ML</u>	<u>A213761</u>	<u>001</u>	Jun 02, 2021
<u>AA</u>		<u>100MG/5ML</u>	<u>A213761</u>	<u>002</u>	Jun 02, 2021
<u>AA</u>	SPECGX LLC	<u>5MG/5ML</u>	<u>A210758</u>	<u>001</u>	Apr 30, 2018
<u>AA</u>		<u>100MG/5ML</u>	<u>A210758</u>	<u>002</u>	Apr 30, 2018
<u>AA</u>	WES PHARMA INC	<u>5MG/5ML</u>	<u>A207511</u>	<u>001</u>	Nov 23, 2016
<u>AA</u>		<u>100MG/5ML</u>	<u>A209897</u>	<u>001</u>	Sep 06, 2017

TABLET;ORAL

OXYCODONE HYDROCHLORIDE

<u>AB</u>	ALVOGEN	<u>5MG</u>	<u>A202116</u>	<u>001</u>	Dec 30, 2011
<u>AB</u>		<u>15MG</u>	<u>A202116</u>	<u>002</u>	Dec 30, 2011
<u>AB</u>		<u>30MG</u>	<u>A202116</u>	<u>003</u>	Dec 30, 2011
<u>AB</u>	AMNEAL PHARMS	<u>5MG</u>	<u>A203638</u>	<u>001</u>	Jun 03, 2014
<u>AB</u>		<u>10MG</u>	<u>A203638</u>	<u>002</u>	Jun 03, 2014
<u>AB</u>		<u>15MG</u>	<u>A203638</u>	<u>003</u>	Jun 03, 2014
<u>AB</u>		<u>20MG</u>	<u>A203638</u>	<u>004</u>	Jun 03, 2014

PRESCRIPTION DRUG PRODUCT LIST

OXYCODONE HYDROCHLORIDE

TABLET; ORAL

OXYCODONE HYDROCHLORIDE

<u>AB</u>		<u>30MG</u>	<u>A203638</u>	<u>005</u>	Jun 03, 2014
<u>AB</u>	ASCENT PHARMS INC	<u>15MG</u>	<u>A207418</u>	<u>001</u>	Aug 07, 2017
<u>AB</u>		<u>30MG</u>	<u>A207418</u>	<u>002</u>	Aug 07, 2017
<u>AB</u>	AUROLIFE PHARMA LLC	<u>5MG</u>	<u>A202160</u>	<u>001</u>	Nov 19, 2012
<u>AB</u>		<u>15MG</u>	<u>A202160</u>	<u>002</u>	Nov 19, 2012
<u>AB</u>		<u>30MG</u>	<u>A202160</u>	<u>003</u>	Nov 19, 2012
<u>AB</u>	AVANTHI INC	<u>5MG</u>	<u>A091393</u>	<u>001</u>	Aug 31, 2009
<u>AB</u>	!	<u>10MG</u>	<u>A091393</u>	<u>002</u>	Aug 31, 2009
<u>AB</u>		<u>15MG</u>	<u>A091393</u>	<u>003</u>	Aug 31, 2009
<u>AB</u>		<u>20MG</u>	<u>A091393</u>	<u>004</u>	Aug 31, 2009
<u>AB</u>		<u>30MG</u>	<u>A091393</u>	<u>005</u>	Aug 31, 2009
<u>AB</u>	EPIC PHARMA LLC	<u>5MG</u>	<u>A090895</u>	<u>001</u>	Aug 24, 2009
<u>AB</u>		<u>5MG</u>	<u>A202662</u>	<u>001</u>	Sep 22, 2015
<u>AB</u>		<u>10MG</u>	<u>A202662</u>	<u>002</u>	Sep 22, 2015
<u>AB</u>		<u>15MG</u>	<u>A090895</u>	<u>002</u>	Aug 24, 2009
<u>AB</u>		<u>15MG</u>	<u>A202662</u>	<u>003</u>	Sep 22, 2015
<u>AB</u>		<u>20MG</u>	<u>A202662</u>	<u>005</u>	Apr 27, 2017
<u>AB</u>		<u>30MG</u>	<u>A090895</u>	<u>003</u>	Aug 24, 2009
<u>AB</u>		<u>30MG</u>	<u>A202662</u>	<u>004</u>	Sep 22, 2015
<u>AB</u>	NOVEL LABS INC	<u>5MG</u>	<u>A204021</u>	<u>001</u>	Jun 12, 2017
<u>AB</u>		<u>10MG</u>	<u>A204021</u>	<u>002</u>	Jun 12, 2017
<u>AB</u>		<u>15MG</u>	<u>A204021</u>	<u>003</u>	Jun 12, 2017
<u>AB</u>		<u>20MG</u>	<u>A204021</u>	<u>004</u>	Jun 12, 2017
<u>AB</u>		<u>30MG</u>	<u>A204021</u>	<u>005</u>	Jun 12, 2017
<u>AB</u>	NUVO PHARM	<u>5MG</u>	<u>A207119</u>	<u>001</u>	Apr 12, 2016
<u>AB</u>		<u>10MG</u>	<u>A207119</u>	<u>002</u>	Apr 12, 2016
<u>AB</u>		<u>15MG</u>	<u>A207119</u>	<u>003</u>	Apr 12, 2016
<u>AB</u>		<u>20MG</u>	<u>A207119</u>	<u>004</u>	Apr 12, 2016
<u>AB</u>		<u>30MG</u>	<u>A207119</u>	<u>005</u>	Apr 12, 2016
<u>AB</u>	RHODES PHARMS	<u>5MG</u>	<u>A091490</u>	<u>001</u>	Mar 09, 2011
<u>AB</u>		<u>10MG</u>	<u>A091490</u>	<u>002</u>	Mar 09, 2011
<u>AB</u>		<u>15MG</u>	<u>A091490</u>	<u>003</u>	Mar 09, 2011
<u>AB</u>		<u>20MG</u>	<u>A091490</u>	<u>004</u>	Mar 09, 2011
<u>AB</u>		<u>30MG</u>	<u>A091490</u>	<u>005</u>	Mar 09, 2011
<u>AB</u>	SPECGX LLC	<u>5MG</u>	<u>A076758</u>	<u>003</u>	Mar 19, 2007
<u>AB</u>		<u>10MG</u>	<u>A076758</u>	<u>004</u>	Oct 12, 2021
<u>AB</u>		<u>15MG</u>	<u>A076758</u>	<u>001</u>	Jun 30, 2004
<u>AB</u>		<u>20MG</u>	<u>A076758</u>	<u>005</u>	Oct 12, 2021
<u>AB</u>		<u>30MG</u>	<u>A076758</u>	<u>002</u>	Jun 30, 2004
<u>AB</u>	STRIDES PHARMA	<u>5MG</u>	<u>A077712</u>	<u>003</u>	Mar 02, 2009
<u>AB</u>		<u>10MG</u>	<u>A077712</u>	<u>004</u>	Apr 13, 2015
<u>AB</u>		<u>15MG</u>	<u>A077712</u>	<u>001</u>	Jan 31, 2007
<u>AB</u>		<u>20MG</u>	<u>A077712</u>	<u>005</u>	Apr 13, 2015
<u>AB</u>		<u>30MG</u>	<u>A077712</u>	<u>002</u>	Jan 31, 2007

ROXICODONE

<u>AB</u>	+	SPECGX LLC	<u>5MG</u>	<u>N021011</u>	<u>003</u>	May 15, 2009
<u>AB</u>	+	!	<u>15MG</u>	<u>N021011</u>	<u>001</u>	Aug 31, 2000
<u>AB</u>	+		<u>30MG</u>	<u>N021011</u>	<u>002</u>	Aug 31, 2000

ROXYBOND

PROTEGA PHARMS	5MG	N209777	001	Apr 20, 2017
	10MG	N209777	004	Sep 05, 2024
	15MG	N209777	002	Apr 20, 2017
	30MG	N209777	003	Apr 20, 2017

TABLET, EXTENDED RELEASE; ORAL

OXYCONTIN

+	PURDUE PHARMA LP	10MG	N022272	001	Apr 05, 2010
+		15MG	N022272	002	Apr 05, 2010
+		20MG	N022272	003	Apr 05, 2010
+		30MG	N022272	004	Apr 05, 2010
+	!	40MG	N022272	005	Apr 05, 2010
+		60MG	N022272	006	Apr 05, 2010
+		80MG	N022272	007	Apr 05, 2010

OXYMETAZOLINE HYDROCHLORIDE

CREAM; TOPICAL

RHOFADE

+	MAYNE PHARMA	1%	N208552	001	Jan 18, 2017
---	--------------	----	---------	-----	--------------

PRESCRIPTION DRUG PRODUCT LISTOXYMETAZOLINE HYDROCHLORIDE

SOLUTION/DROPS;OPHTHALMIC

UPNEEQ

+! RVL PHARMS 0.1% N212520 001 Jul 08, 2020

OXYMETAZOLINE HYDROCHLORIDE; TETRACAINE HYDROCHLORIDE

SPRAY, METERED;NASAL

KOVANAZE

+! ST RENATUS 0.1MG/SPRAY;6MG/SPRAY N208032 001 Jun 29, 2016

OXYMORPHONE HYDROCHLORIDE

TABLET;ORAL

OXYMORPHONE HYDROCHLORIDE

<u>AB</u>	ASCENT PHARMS INC	<u>5MG</u>	<u>A210175</u>	<u>001</u>	Feb 02, 2018
<u>AB</u>		<u>10MG</u>	<u>A210175</u>	<u>002</u>	Feb 02, 2018
<u>AB</u>	AUROLIFE PHARMA LLC	<u>5MG</u>	<u>A204459</u>	<u>001</u>	Apr 26, 2016
<u>AB</u>		<u>10MG</u>	<u>A204459</u>	<u>002</u>	Apr 26, 2016
<u>AB</u>	AVANTHI INC	<u>5MG</u>	<u>A203601</u>	<u>001</u>	Jan 30, 2013
<u>AB</u>	!	<u>10MG</u>	<u>A203601</u>	<u>002</u>	Jan 30, 2013
<u>AB</u>	EPIC PHARMA LLC	<u>5MG</u>	<u>A201187</u>	<u>001</u>	Dec 15, 2014
<u>AB</u>		<u>10MG</u>	<u>A201187</u>	<u>002</u>	Dec 15, 2014
<u>AB</u>	HIKMA	<u>5MG</u>	<u>A090964</u>	<u>001</u>	Sep 27, 2010
<u>AB</u>		<u>10MG</u>	<u>A090964</u>	<u>002</u>	Sep 27, 2010
<u>AB</u>	TEVA	<u>5MG</u>	<u>A091443</u>	<u>002</u>	Feb 15, 2011
<u>AB</u>		<u>10MG</u>	<u>A091443</u>	<u>001</u>	Feb 15, 2011

TABLET, EXTENDED RELEASE;ORAL

OXYMORPHONE HYDROCHLORIDE

IMPAX LABS

	5MG	A079087	001	Jun 14, 2010
	7.5MG	A079087	002	Dec 21, 2010
	10MG	A079087	003	Jun 14, 2010
	15MG	A079087	004	Dec 21, 2010
	20MG	A079087	005	Jun 14, 2010
	30MG	A079087	006	Jul 22, 2010
!	40MG	A079087	007	Jun 14, 2010

OXYTOCIN

INJECTABLE;INJECTION

OXYTOCIN

<u>AP</u>	+!	FRESENIUS KABI USA	<u>10USP UNITS/ML (10USP UNITS/ML)</u>	<u>N018248</u>	<u>001</u>	
<u>AP</u>	+		<u>100USP UNITS/10ML (10USP UNITS/ML)</u>	<u>N018248</u>	<u>002</u>	
<u>AP</u>	+	HIKMA	<u>10USP UNITS/ML (10USP UNITS/ML)</u>	<u>N018243</u>	<u>001</u>	
<u>AP</u>	+		<u>100USP UNITS/10ML (10USP UNITS/ML)</u>	<u>N018243</u>	<u>002</u>	Jan 10, 2007
<u>AP</u>		HIKMA FARMACEUTICA	<u>10USP UNITS/ML (10USP UNITS/ML)</u>	<u>A200219</u>	<u>001</u>	Feb 13, 2013
<u>AP</u>		SAGENT PHARMS INC	<u>10USP UNITS/ML (10USP UNITS/ML)</u>	<u>A091676</u>	<u>001</u>	Jul 13, 2018
<u>AP</u>			<u>100USP UNITS/10ML (10USP UNITS/ML)</u>	<u>A091676</u>	<u>002</u>	Jul 13, 2018

PITOCIN

<u>AP</u>	+!	ENDO OPERATIONS	<u>10USP UNITS/ML (10USP UNITS/ML)</u>	<u>N018261</u>	<u>001</u>	
<u>AP</u>	+		<u>100USP UNITS/10ML (10USP UNITS/ML)</u>	<u>N018261</u>	<u>002</u>	Jul 27, 2007
OXYTOCIN						
	+!	FRESENIUS KABI USA	300USP UNITS/30ML (10USP UNITS/ML)	N018248	003	Jul 27, 2007
PITOCIN						
	+	ENDO OPERATIONS	500USP UNITS/50ML (10USP UNITS/ML)	N018261	003	Sep 05, 2012

OZANIMOD HYDROCHLORIDE

CAPSULE;ORAL

ZEPOSIA

+	BRISTOL	EQ 0.23MG BASE	N209899	001	Mar 25, 2020
+		EQ 0.46MG BASE	N209899	002	Mar 25, 2020
+!		EQ 0.92MG BASE	N209899	003	Mar 25, 2020

PACLITAXEL

INJECTABLE;INJECTION

PACLITAXEL

<u>AP</u>		ACCORD HLTHCARE	<u>6MG/ML</u>	<u>A205720</u>	<u>001</u>	Aug 17, 2018
<u>AP</u>		ACTAVIS TOTOWA	<u>6MG/ML</u>	<u>A090130</u>	<u>001</u>	Dec 09, 2009
<u>AP</u>		ALEMBIC	<u>6MG/ML</u>	<u>A216874</u>	<u>001</u>	Oct 20, 2022
<u>AP</u>		FRESENIUS KABI USA	<u>6MG/ML</u>	<u>A077574</u>	<u>001</u>	Nov 27, 2006
<u>AP</u>		GLAND PHARMA LTD	<u>6MG/ML</u>	<u>A207326</u>	<u>001</u>	Aug 23, 2016
<u>AP</u>		HIKMA	<u>6MG/ML</u>	<u>A075190</u>	<u>001</u>	Jan 28, 2002
<u>AP</u>	!	HOSPIRA	<u>6MG/ML</u>	<u>A076131</u>	<u>001</u>	May 08, 2002
<u>AP</u>		MSN	<u>6MG/ML</u>	<u>A213434</u>	<u>001</u>	Aug 24, 2020
<u>AP</u>		TEVA PHARMS	<u>6MG/ML</u>	<u>A075184</u>	<u>001</u>	Jan 25, 2002

PRESCRIPTION DRUG PRODUCT LIST

PACLITAXEL

POWDER; INTRAVENOUS

ABRAXANE

<u>AB</u>	+ !	BRISTOL-MYERS	<u>100MG/VIAL</u>	<u>N021660</u>	<u>001</u>	Jan 07, 2005
-----------	------------	---------------	-------------------	----------------	------------	--------------

PACLITAXEL

<u>AB</u>	+ !	AM REGENT	<u>100MG/VIAL</u>	<u>N211875</u>	<u>001</u>	Jul 27, 2022
-----------	------------	-----------	-------------------	----------------	------------	--------------

<u>AB</u>		HENGRUI PHARMA	<u>100MG/VIAL</u>	<u>A212700</u>	<u>001</u>	Oct 08, 2024
-----------	--	----------------	-------------------	----------------	------------	--------------

<u>AB</u>	+ !	TEVA PHARMS INC	<u>100MG/VIAL</u>	<u>N216338</u>	<u>001</u>	May 11, 2023
-----------	------------	-----------------	-------------------	----------------	------------	--------------

PACRITINIB CITRATE

CAPSULE; ORAL

VONJO

+ !	SOBI	EQ 100MG BASE	N208712	001	Feb 28, 2022
------------	------	---------------	---------	-----	--------------

PAFOLACIANINE SODIUM

SOLUTION; INTRAVENOUS

CYTALUX

+ !	ON TARGET LABS	EQ 3.2MG BASE/1.6ML (EQ 2MG BASE/ML)	N214907	001	Nov 29, 2021
------------	----------------	--------------------------------------	---------	-----	--------------

PALBOCICLIB

CAPSULE; ORAL

IBRANCE

+	PFIZER	75MG	N207103	001	Feb 03, 2015
----------	--------	------	---------	-----	--------------

+		100MG	N207103	002	Feb 03, 2015
----------	--	-------	---------	-----	--------------

+ !		125MG	N207103	003	Feb 03, 2015
------------	--	-------	---------	-----	--------------

TABLET; ORAL

IBRANCE

+	PFIZER	75MG	N212436	001	Nov 01, 2019
----------	--------	------	---------	-----	--------------

+		100MG	N212436	002	Nov 01, 2019
----------	--	-------	---------	-----	--------------

+ !		125MG	N212436	003	Nov 01, 2019
------------	--	-------	---------	-----	--------------

PALIPERIDONE

TABLET, EXTENDED RELEASE; ORAL

INVEGA

<u>AB</u>	+	JANSSEN PHARMS	<u>3MG</u>	<u>N021999</u>	<u>001</u>	Dec 19, 2006
-----------	----------	----------------	------------	----------------	------------	--------------

<u>AB</u>	+ !		<u>6MG</u>	<u>N021999</u>	<u>002</u>	Dec 19, 2006
-----------	------------	--	------------	----------------	------------	--------------

<u>AB</u>	+		<u>9MG</u>	<u>N021999</u>	<u>003</u>	Dec 19, 2006
-----------	----------	--	------------	----------------	------------	--------------

PALIPERIDONE

<u>AB</u>		AJANTA PHARMA LTD	<u>1.5MG</u>	<u>A218514</u>	<u>001</u>	Jun 26, 2024
-----------	--	-------------------	--------------	----------------	------------	--------------

<u>AB</u>		ALEMBIC	<u>1.5MG</u>	<u>A218330</u>	<u>001</u>	Sep 26, 2024
-----------	--	---------	--------------	----------------	------------	--------------

<u>AB</u>			<u>3MG</u>	<u>A218330</u>	<u>002</u>	Sep 26, 2024
-----------	--	--	------------	----------------	------------	--------------

<u>AB</u>			<u>6MG</u>	<u>A218330</u>	<u>003</u>	Sep 26, 2024
-----------	--	--	------------	----------------	------------	--------------

<u>AB</u>			<u>9MG</u>	<u>A218330</u>	<u>004</u>	Sep 26, 2024
-----------	--	--	------------	----------------	------------	--------------

<u>AB</u>		AMNEAL PHARMS	<u>1.5MG</u>	<u>A204707</u>	<u>001</u>	Sep 23, 2019
-----------	--	---------------	--------------	----------------	------------	--------------

<u>AB</u>			<u>3MG</u>	<u>A204707</u>	<u>002</u>	Sep 23, 2019
-----------	--	--	------------	----------------	------------	--------------

<u>AB</u>			<u>6MG</u>	<u>A204707</u>	<u>003</u>	Sep 23, 2019
-----------	--	--	------------	----------------	------------	--------------

<u>AB</u>			<u>9MG</u>	<u>A204707</u>	<u>004</u>	Sep 23, 2019
-----------	--	--	------------	----------------	------------	--------------

<u>AB</u>		ASCENT PHARMS INC	<u>1.5MG</u>	<u>A216174</u>	<u>001</u>	Aug 23, 2023
-----------	--	-------------------	--------------	----------------	------------	--------------

<u>AB</u>			<u>3MG</u>	<u>A216174</u>	<u>002</u>	Aug 23, 2023
-----------	--	--	------------	----------------	------------	--------------

<u>AB</u>			<u>6MG</u>	<u>A216174</u>	<u>003</u>	Aug 23, 2023
-----------	--	--	------------	----------------	------------	--------------

<u>AB</u>			<u>9MG</u>	<u>A216174</u>	<u>004</u>	Aug 23, 2023
-----------	--	--	------------	----------------	------------	--------------

<u>AB</u>		CSPC OUYI	<u>1.5MG</u>	<u>A212807</u>	<u>001</u>	Oct 29, 2020
-----------	--	-----------	--------------	----------------	------------	--------------

<u>AB</u>			<u>3MG</u>	<u>A212807</u>	<u>002</u>	Oct 29, 2020
-----------	--	--	------------	----------------	------------	--------------

<u>AB</u>			<u>6MG</u>	<u>A212807</u>	<u>003</u>	Oct 29, 2020
-----------	--	--	------------	----------------	------------	--------------

<u>AB</u>			<u>9MG</u>	<u>A212807</u>	<u>004</u>	Oct 29, 2020
-----------	--	--	------------	----------------	------------	--------------

<u>AB</u>		INVENTIA	<u>1.5MG</u>	<u>A204452</u>	<u>001</u>	Jun 12, 2019
-----------	--	----------	--------------	----------------	------------	--------------

<u>AB</u>			<u>3MG</u>	<u>A204452</u>	<u>002</u>	Jun 12, 2019
-----------	--	--	------------	----------------	------------	--------------

<u>AB</u>			<u>6MG</u>	<u>A204452</u>	<u>003</u>	Jun 12, 2019
-----------	--	--	------------	----------------	------------	--------------

<u>AB</u>			<u>9MG</u>	<u>A204452</u>	<u>004</u>	Jun 12, 2019
-----------	--	--	------------	----------------	------------	--------------

<u>AB</u>		RK PHARMA	<u>1.5MG</u>	<u>A203802</u>	<u>001</u>	Sep 24, 2015
-----------	--	-----------	--------------	----------------	------------	--------------

<u>AB</u>			<u>3MG</u>	<u>A203802</u>	<u>002</u>	Sep 24, 2015
-----------	--	--	------------	----------------	------------	--------------

<u>AB</u>			<u>6MG</u>	<u>A203802</u>	<u>003</u>	Sep 24, 2015
-----------	--	--	------------	----------------	------------	--------------

<u>AB</u>			<u>9MG</u>	<u>A203802</u>	<u>004</u>	Sep 24, 2015
-----------	--	--	------------	----------------	------------	--------------

<u>AB</u>		SUN PHARM	<u>1.5MG</u>	<u>A205618</u>	<u>001</u>	Apr 06, 2018
-----------	--	-----------	--------------	----------------	------------	--------------

<u>AB</u>			<u>3MG</u>	<u>A205618</u>	<u>002</u>	Apr 06, 2018
-----------	--	--	------------	----------------	------------	--------------

<u>AB</u>			<u>6MG</u>	<u>A205618</u>	<u>003</u>	Apr 06, 2018
-----------	--	--	------------	----------------	------------	--------------

<u>AB</u>			<u>9MG</u>	<u>A205618</u>	<u>004</u>	Apr 06, 2018
-----------	--	--	------------	----------------	------------	--------------

<u>AB</u>		ZYDUS PHARMS	<u>1.5MG</u>	<u>A217445</u>	<u>001</u>	Oct 08, 2024
-----------	--	--------------	--------------	----------------	------------	--------------

<u>AB</u>			<u>3MG</u>	<u>A217445</u>	<u>002</u>	Oct 08, 2024
-----------	--	--	------------	----------------	------------	--------------

<u>AB</u>			<u>6MG</u>	<u>A217445</u>	<u>003</u>	Oct 08, 2024
-----------	--	--	------------	----------------	------------	--------------

<u>AB</u>			<u>9MG</u>	<u>A217445</u>	<u>004</u>	Oct 08, 2024
-----------	--	--	------------	----------------	------------	--------------

BX		LUPIN LTD	1.5MG	A208643	001	Jun 29, 2022
----	--	-----------	-------	---------	-----	--------------

BX			3MG	A208643	002	Jun 29, 2022
----	--	--	-----	---------	-----	--------------

BX			6MG	A208643	003	Jun 29, 2022
----	--	--	-----	---------	-----	--------------

PRESCRIPTION DRUG PRODUCT LIST

PALIPERIDONE

TABLET, EXTENDED RELEASE;ORAL

PALIPERIDONE

BX		9MG		A208643	004	Jun 29, 2022
----	--	-----	--	---------	-----	--------------

PALIPERIDONE PALMITATE

SUSPENSION, EXTENDED RELEASE;INTRAMUSCULAR

INVEGA SUSTENNA

<u>AB</u>	+	JANSSEN PHARMS	<u>39MG/0.25ML (39MG/0.25ML)</u>	<u>N022264</u>	<u>001</u>	Jul 31, 2009
<u>AB</u>	+		<u>78MG/0.5ML (78MG/0.5ML)</u>	<u>N022264</u>	<u>002</u>	Jul 31, 2009
<u>AB</u>	+		<u>117MG/0.75ML (117MG/0.75ML)</u>	<u>N022264</u>	<u>003</u>	Jul 31, 2009
<u>AB</u>	+		<u>156MG/ML (156MG/ML)</u>	<u>N022264</u>	<u>004</u>	Jul 31, 2009
<u>AB</u>	+		<u>234MG/1.5ML (156MG/ML)</u>	<u>N022264</u>	<u>005</u>	Jul 31, 2009

PALIPERIDONE PALMITATE

<u>AB</u>		TEVA PHARMS USA	<u>39MG/0.25ML (39MG/0.25ML)</u>	<u>A211149</u>	<u>001</u>	Jul 06, 2021
<u>AB</u>			<u>78MG/0.5ML (78MG/0.5ML)</u>	<u>A211149</u>	<u>002</u>	Jul 06, 2021
<u>AB</u>			<u>117MG/0.75ML (117MG/0.75ML)</u>	<u>A211149</u>	<u>003</u>	Jul 06, 2021
<u>AB</u>			<u>156MG/ML (156MG/ML)</u>	<u>A211149</u>	<u>004</u>	Jul 06, 2021
<u>AB</u>			<u>234MG/1.5ML (156MG/ML)</u>	<u>A211149</u>	<u>005</u>	Jul 06, 2021

ERZOFRI

+	!	LUYE INNOMIND PHARMA	39MG/0.25ML (39MG/0.25ML)	N216352	001	Jul 26, 2024
+	!		78MG/0.5ML (78MG/0.5ML)	N216352	002	Jul 26, 2024
+	!		117MG/0.75ML (117MG/0.75ML)	N216352	003	Jul 26, 2024
+	!		156MG/ML (156MG/ML)	N216352	004	Jul 26, 2024
+	!		234MG/1.5ML (156MG/ML)	N216352	005	Jul 26, 2024
+	!		351MG/2.25ML (156MG/ML)	N216352	006	Jul 26, 2024

INVEGA HAFYERA

+		JANSSEN PHARMS	1.092GM/3.5ML (312MG/ML)	N207946	005	Aug 30, 2021
+			1.560GM/5ML (312MG/ML)	N207946	006	Aug 30, 2021

INVEGA TRINZA

+	!	JANSSEN PHARMS	273MG/0.875ML (273MG/0.875ML)	N207946	001	May 18, 2015
+	!		410MG/1.315ML (311.79MG/ML)	N207946	002	May 18, 2015
+	!		546MG/1.75ML (312MG/ML)	N207946	003	May 18, 2015
+	!		819MG/2.625ML (312MG/ML)	N207946	004	May 18, 2015

PALONOSETRON HYDROCHLORIDE

INJECTABLE; INTRAVENOUS

PALONOSETRON HYDROCHLORIDE

<u>AP</u>		AVET LIFESCIENCES	<u>EQ 0.25MG BASE/5ML (EQ 0.05MG BASE/ML)</u>	<u>A202951</u>	<u>002</u>	Jun 29, 2021
<u>AP</u>		BAXTER HLTHCARE CORP	<u>EQ 0.25MG BASE/5ML (EQ 0.05MG BASE/ML)</u>	<u>A206916</u>	<u>001</u>	Nov 12, 2021
<u>AP</u>		CHARTWELL RX	<u>EQ 0.25MG BASE/5ML (EQ 0.05MG BASE/ML)</u>	<u>A209287</u>	<u>001</u>	Sep 19, 2018
<u>AP</u>	!	DR REDDYS	<u>EQ 0.25MG BASE/5ML (EQ 0.05MG BASE/ML)</u>	<u>A201533</u>	<u>002</u>	Apr 21, 2016
<u>AP</u>		EUGIA PHARMA	<u>EQ 0.25MG BASE/5ML (EQ 0.05MG BASE/ML)</u>	<u>A204702</u>	<u>001</u>	Nov 06, 2018
<u>AP</u>		FRESENIUS KABI USA	<u>EQ 0.25MG BASE/5ML (EQ 0.05MG BASE/ML)</u>	<u>A206801</u>	<u>001</u>	Sep 19, 2018
<u>AP</u>			<u>EQ 0.25MG BASE/5ML (EQ 0.05MG BASE/ML)</u>	<u>A206802</u>	<u>001</u>	Sep 19, 2018
<u>AP</u>		HOSPIRA	<u>EQ 0.25MG BASE/5ML (EQ 0.05MG BASE/ML)</u>	<u>A207005</u>	<u>001</u>	Sep 19, 2018
<u>AP</u>		MEITHEAL	<u>EQ 0.25MG BASE/5ML (EQ 0.05MG BASE/ML)</u>	<u>A215861</u>	<u>001</u>	Aug 14, 2023
<u>AP</u>		MYLAN INSTITUTIONAL	<u>EQ 0.25MG BASE/5ML (EQ 0.05MG BASE/ML)</u>	<u>A206416</u>	<u>001</u>	Sep 19, 2018
<u>AP</u>		QILU PHARM HAINAN	<u>EQ 0.25MG BASE/5ML (EQ 0.05MG BASE/ML)</u>	<u>A205648</u>	<u>001</u>	Sep 19, 2018
<u>AP</u>		SAGENT PHARMS INC	<u>EQ 0.25MG BASE/5ML (EQ 0.05MG BASE/ML)</u>	<u>A204289</u>	<u>001</u>	Sep 19, 2018
<u>AP</u>			<u>EQ 0.25MG BASE/5ML (EQ 0.05MG BASE/ML)</u>	<u>A205870</u>	<u>001</u>	Sep 19, 2018
<u>AP</u>		SANDOZ	<u>EQ 0.25MG BASE/5ML (EQ 0.05MG BASE/ML)</u>	<u>A202521</u>	<u>001</u>	Oct 13, 2015
<u>AP</u>		TEVA PHARMS USA	<u>EQ 0.25MG BASE/5ML (EQ 0.05MG BASE/ML)</u>	<u>A090713</u>	<u>001</u>	Mar 23, 2018
	!	AVET LIFESCIENCES	EQ 0.075MG BASE/1.5ML (EQ 0.05MG BASE/ML)	A202951	001	Jun 29, 2021

SOLUTION; INTRAVENOUS

POSFREA

+	!	AVYXA HOLDINGS	EQ 0.25MG BASE/5ML (EQ 0.05MG BASE/ML)	N203050	002	Mar 01, 2016
---	---	----------------	--	---------	-----	--------------

PALOPEGTERIPARATIDE

SOLUTION; SUBCUTANEOUS

YORVIPATH

+	!	ASCENDIS PHARMA BONE	EQ 0.168MG TERIPARATIDE/0.56ML (EQ 0.168MG TERIPARATIDE/0.56ML)	N216490	001	Aug 09, 2024
+			EQ 0.294MG TERIPARATIDE/0.98ML (EQ 0.294MG TERIPARATIDE/0.98ML)	N216490	002	Aug 09, 2024
+			EQ 0.42MG TERIPARATIDE/1.4ML (EQ 0.3MG TERIPARATIDE/ML)	N216490	003	Aug 09, 2024

PRESCRIPTION DRUG PRODUCT LIST

PALOVAROTENE

CAPSULE; ORAL

SOHONOS

+	IPSEN	1MG	N215559	001	Aug 16, 2023
+		1.5MG	N215559	002	Aug 16, 2023
+		2.5MG	N215559	003	Aug 16, 2023
+		5MG	N215559	004	Aug 16, 2023
+	!	10MG	N215559	005	Aug 16, 2023

PAMIDRONATE DISODIUM

INJECTABLE; INJECTION

PAMIDRONATE DISODIUM

<u>AP</u>	AREVA PHARMS	<u>30MG/VIAL</u>	<u>A077433</u>	<u>001</u>	Nov 26, 2008
<u>AP</u>		<u>90MG/VIAL</u>	<u>A077433</u>	<u>003</u>	Nov 26, 2008
<u>AP</u>	DR REDDYS	<u>60MG/10ML (6MG/ML)</u>	<u>A078156</u>	<u>002</u>	Aug 19, 2008
<u>AP</u>	HIKMA	<u>30MG/VIAL</u>	<u>A075290</u>	<u>001</u>	Apr 30, 2001
<u>AP</u>		<u>90MG/VIAL</u>	<u>A075290</u>	<u>003</u>	Apr 30, 2001
<u>AP</u>	!	<u>60MG/10ML (6MG/ML)</u>	<u>A075841</u>	<u>002</u>	Jun 27, 2002
<u>AP1</u>	+	<u>30MG/10ML (3MG/ML)</u>	<u>N021113</u>	<u>001</u>	Mar 04, 2002
<u>AP1</u>	+	<u>90MG/10ML (9MG/ML)</u>	<u>N021113</u>	<u>002</u>	Mar 04, 2002
<u>AP1</u>	SAGENT PHARMS INC	<u>30MG/10ML (3MG/ML)</u>	<u>A078373</u>	<u>001</u>	Dec 23, 2008
<u>AP1</u>		<u>90MG/10ML (9MG/ML)</u>	<u>A078373</u>	<u>002</u>	Dec 23, 2008
<u>AP2</u>	DR REDDYS	<u>30MG/10ML (3MG/ML)</u>	<u>A078156</u>	<u>001</u>	Aug 19, 2008
<u>AP2</u>		<u>90MG/10ML (9MG/ML)</u>	<u>A078156</u>	<u>003</u>	Aug 19, 2008
<u>AP2</u>	HOSPIRA	<u>30MG/10ML (3MG/ML)</u>	<u>A075841</u>	<u>001</u>	Jun 27, 2002
<u>AP2</u>		<u>90MG/10ML (9MG/ML)</u>	<u>A075841</u>	<u>003</u>	Jun 27, 2002
<u>AP2</u>	MYLAN LABS LTD	<u>30MG/10ML (3MG/ML)</u>	<u>A078520</u>	<u>001</u>	Oct 31, 2008
<u>AP2</u>		<u>90MG/10ML (9MG/ML)</u>	<u>A078520</u>	<u>002</u>	Oct 31, 2008
	AREVA PHARMS	60MG/VIAL	A077433	002	Nov 26, 2008

PANCURONIUM BROMIDE

INJECTABLE; INJECTION

PANCURONIUM BROMIDE

!	DR REDDYS	1MG/ML	A072759	001	Jul 31, 1990
!		2MG/ML	A072760	001	Jul 31, 1990

PANTOPRAZOLE SODIUM

FOR SUSPENSION, DELAYED RELEASE; ORAL

PANTOPRAZOLE SODIUM

<u>AB</u>	AJANTA PHARMA LTD	<u>EQ 40MG BASE</u>	<u>A217416</u>	<u>001</u>	Feb 09, 2023
<u>AB</u>	ANNORA PHARMA	<u>EQ 40MG BASE</u>	<u>A216139</u>	<u>001</u>	Oct 27, 2023
<u>AB</u>	DEXCEL	<u>EQ 40MG BASE</u>	<u>A216247</u>	<u>001</u>	Jun 16, 2023
<u>AB</u>	SUN PHARM	<u>EQ 40MG BASE</u>	<u>A213725</u>	<u>001</u>	Jun 30, 2020

PROTONIX

<u>AB</u>	+	WYETH PHARMS	<u>EQ 40MG BASE</u>	<u>N022020</u>	<u>001</u>	Nov 14, 2007
-----------	---	--------------	---------------------	----------------	------------	--------------

INJECTABLE; INTRAVENOUS

PANTOPRAZOLE SODIUM

<u>AP</u>	ASPIRO	<u>EQ 40MG BASE/VIAL</u>	<u>A213778</u>	<u>001</u>	May 18, 2022
<u>AP</u>	BE PHARMS	<u>EQ 40MG BASE/VIAL</u>	<u>A216171</u>	<u>001</u>	May 18, 2022
<u>AP</u>	EPIC PHARMA LLC	<u>EQ 40MG BASE/VIAL</u>	<u>A079197</u>	<u>001</u>	Nov 08, 2012
<u>AP</u>	EUGIA PHARMA	<u>EQ 40MG BASE/VIAL</u>	<u>A205675</u>	<u>001</u>	Mar 30, 2016
<u>AP</u>	GLAND PHARMA LTD	<u>EQ 40MG BASE/VIAL</u>	<u>A204400</u>	<u>001</u>	May 18, 2022
<u>AP</u>	HANGZHOU ZHONGMEI	<u>EQ 40MG BASE/VIAL</u>	<u>A209524</u>	<u>001</u>	Aug 30, 2021
<u>AP</u>	KNACK	<u>EQ 40MG BASE/VIAL</u>	<u>A214680</u>	<u>001</u>	May 19, 2022
<u>AP</u>	MEITHEAL	<u>EQ 40MG BASE/VIAL</u>	<u>A215860</u>	<u>001</u>	Aug 29, 2022

PROTONIX IV

<u>AP</u>	+	WYETH PHARMS	<u>EQ 40MG BASE/VIAL</u>	<u>N020988</u>	<u>001</u>	Mar 22, 2001
-----------	---	--------------	--------------------------	----------------	------------	--------------

POWDER; INTRAVENOUS

PANTOPRAZOLE SODIUM

<u>AP</u>	FRESENIUS KABI USA	<u>EQ 40MG BASE/VIAL</u>	<u>A216021</u>	<u>001</u>	Aug 01, 2024	
<u>AP</u>	+	HIKMA	<u>EQ 40MG BASE/VIAL</u>	<u>N209463</u>	<u>001</u>	Jun 30, 2017

SOLUTION; INTRAVENOUS

PANTOPRAZOLE SODIUM IN 0.9% SODIUM CHLORIDE

+	BAXTER HLTHCARE CORP	EQ 40MG BASE/100ML (EQ 0.4MG BASE/ML)	N217512	001	Feb 14, 2024
+		EQ 40MG BASE/50ML (EQ 0.8MG BASE/ML)	N217512	002	Feb 14, 2024
+		EQ 80MG BASE/100ML (EQ 0.8MG BASE/ML)	N217512	003	Feb 14, 2024

TABLET, DELAYED RELEASE; ORAL

PANTOPRAZOLE SODIUM

<u>AB</u>	ACTAVIS TOTOWA	<u>EQ 20MG BASE</u>	<u>A090797</u>	<u>001</u>	Feb 07, 2011
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A090797</u>	<u>002</u>	Feb 07, 2011
<u>AB</u>	AMNEAL PHARMS	<u>EQ 20MG BASE</u>	<u>A205119</u>	<u>001</u>	Jan 26, 2016
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A205119</u>	<u>002</u>	Jan 26, 2016
<u>AB</u>	AUROBINDO PHARMA	<u>EQ 20MG BASE</u>	<u>A202038</u>	<u>001</u>	Sep 28, 2012

PRESCRIPTION DRUG PRODUCT LIST

PANTOPRAZOLE SODIUM

TABLET, DELAYED RELEASE;ORAL

PANTOPRAZOLE SODIUM

LTD

<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A202038 002</u>	Sep 28, 2012
<u>AB</u>	DR REDDYS LABS LTD	<u>EQ 20MG BASE</u>	<u>A077619 001</u>	Jan 19, 2011
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A077619 002</u>	Jan 19, 2011
<u>AB</u>	GRANULES	<u>EQ 20MG BASE</u>	<u>A217282 001</u>	Dec 11, 2023
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A217282 002</u>	Dec 11, 2023
<u>AB</u>	HETERO LABS LTD V	<u>EQ 20MG BASE</u>	<u>A202882 001</u>	Sep 10, 2014
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A202882 002</u>	Sep 10, 2014
<u>AB</u>	INGENUS PHARMS LLC	<u>EQ 40MG BASE</u>	<u>A211368 001</u>	Mar 01, 2019
<u>AB</u>	LANNETT CO INC	<u>EQ 20MG BASE</u>	<u>A078281 001</u>	Jan 20, 2011
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A078281 002</u>	Jan 20, 2011
<u>AB</u>	MANKIND PHARMA	<u>EQ 40MG BASE</u>	<u>A215880 001</u>	Jul 26, 2022
<u>AB</u>	MYLAN PHARMS INC	<u>EQ 20MG BASE</u>	<u>A090970 001</u>	Jan 19, 2011
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A090970 002</u>	Jan 19, 2011
<u>AB</u>	ORBION PHARMS	<u>EQ 20MG BASE</u>	<u>A202052 001</u>	Dec 02, 2014
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A202052 002</u>	Dec 02, 2014
<u>AB</u>	RUBICON	<u>EQ 20MG BASE</u>	<u>A090807 001</u>	May 02, 2012
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A090807 002</u>	May 02, 2012
<u>AB</u>	TORRENT PHARMS	<u>EQ 20MG BASE</u>	<u>A090074 001</u>	Jan 19, 2011
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A090074 002</u>	Jan 19, 2011
<u>AB</u>	WOCKHARDT BIO AG	<u>EQ 20MG BASE</u>	<u>A091231 001</u>	Jan 19, 2011
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A091231 002</u>	Jan 19, 2011
<u>PROTONIX</u>				
<u>AB</u>	+ WYETH PHARMS	<u>EQ 20MG BASE</u>	<u>N020987 002</u>	Jun 12, 2001
<u>AB</u>	+!	<u>EQ 40MG BASE</u>	<u>N020987 001</u>	Feb 02, 2000

PARICALCITOL

CAPSULE;ORAL

PARICALCITOL

<u>AB</u>	AMNEAL PHARMS	<u>1MCG</u>	<u>A204327 001</u>	Jan 13, 2016
<u>AB</u>		<u>2MCG</u>	<u>A204327 002</u>	Jan 13, 2016
<u>AB</u>		<u>4MCG</u>	<u>A204327 003</u>	Jan 13, 2016
<u>AB</u>	AUROBINDO PHARMA LTD	<u>1MCG</u>	<u>A207672 001</u>	Jan 14, 2016
<u>AB</u>		<u>2MCG</u>	<u>A207672 002</u>	Jan 14, 2016
<u>AB</u>		<u>4MCG</u>	<u>A207672 003</u>	Jan 14, 2016
<u>AB</u>	BIONPHARMA	<u>1MCG</u>	<u>A202539 001</u>	Mar 27, 2014
<u>AB</u>		<u>2MCG</u>	<u>A202539 002</u>	Mar 27, 2014
<u>AB</u>		<u>4MCG</u>	<u>A202539 003</u>	Mar 27, 2014
<u>AB</u>	DR REDDYS	<u>1MCG</u>	<u>A091412 001</u>	Jun 24, 2014
<u>AB</u>		<u>2MCG</u>	<u>A091412 002</u>	Jun 24, 2014
<u>AB</u>		<u>4MCG</u>	<u>A091412 003</u>	Jun 24, 2014
<u>AB</u>	MARKSANS PHARMA	<u>1MCG</u>	<u>A204948 001</u>	Oct 07, 2016
<u>AB</u>		<u>2MCG</u>	<u>A204948 002</u>	Oct 07, 2016
<u>AB</u>		<u>4MCG</u>	<u>A204948 003</u>	Oct 07, 2016
<u>AB</u>	RISING	<u>1MCG</u>	<u>A202124 001</u>	Jun 24, 2014
<u>AB</u>		<u>2MCG</u>	<u>A202124 002</u>	Jun 24, 2014
<u>AB</u>		<u>4MCG</u>	<u>A202124 003</u>	Jun 24, 2014
<u>AB</u>	TEVA PHARMS USA	<u>1MCG</u>	<u>A090829 001</u>	Sep 27, 2013
<u>AB</u>		<u>2MCG</u>	<u>A090829 002</u>	Sep 27, 2013
<u>AB</u>	!	<u>4MCG</u>	<u>A090829 003</u>	Sep 27, 2013
<u>ZEMPLAR</u>				
<u>AB</u>	+ ABBVIE	<u>1MCG</u>	<u>N021606 001</u>	May 26, 2005
<u>AB</u>	+	<u>2MCG</u>	<u>N021606 002</u>	May 26, 2005

SOLUTION;INTRAVENOUS

PARICALCITOL

<u>AP</u>	ACCORD HLTHCARE	<u>0.002MG/ML (0.002MG/ML)</u>	<u>N207174 001</u>	Feb 04, 2016
<u>AP</u>		<u>0.005MG/ML (0.005MG/ML)</u>	<u>N207174 002</u>	Feb 04, 2016
<u>AP</u>		<u>0.01MG/2ML (0.005MG/ML)</u>	<u>N207174 003</u>	Feb 04, 2016
<u>AP</u>	AMNEAL PHARMS CO	<u>0.002MG/ML (0.002MG/ML)</u>	<u>A206699 001</u>	Mar 09, 2017
<u>AP</u>		<u>0.005MG/ML (0.005MG/ML)</u>	<u>A206699 002</u>	Mar 09, 2017
<u>AP</u>		<u>0.01MG/2ML (0.005MG/ML)</u>	<u>A206699 003</u>	Mar 09, 2017
<u>AP</u>	DR REDDYS	<u>0.002MG/ML (0.002MG/ML)</u>	<u>A204910 001</u>	Aug 17, 2016
<u>AP</u>		<u>0.005MG/ML (0.005MG/ML)</u>	<u>A204910 002</u>	Aug 17, 2016
<u>AP</u>		<u>0.01MG/2ML (0.005MG/ML)</u>	<u>A204910 003</u>	Aug 17, 2016
<u>AP</u>	EUGIA PHARMA	<u>0.002MG/ML (0.002MG/ML)</u>	<u>A205982 001</u>	Oct 09, 2018
<u>AP</u>		<u>0.005MG/ML (0.005MG/ML)</u>	<u>A205982 002</u>	Oct 09, 2018
<u>AP</u>		<u>0.01MG/2ML (0.005MG/ML)</u>	<u>A205982 003</u>	Oct 09, 2018
<u>AP</u>	HIKMA PHARMS	<u>0.002MG/ML (0.002MG/ML)</u>	<u>N205917 001</u>	Nov 18, 2014
<u>AP</u>		<u>0.005MG/ML (0.005MG/ML)</u>	<u>N205917 002</u>	Nov 18, 2014

PRESCRIPTION DRUG PRODUCT LIST

PARICALCITOL

SOLUTION; INTRAVENOUS

PARICALCITOL

<u>AP</u>		<u>0.01MG/2ML (0.005MG/ML)</u>	<u>N205917 003</u>	Nov 18, 2014
<u>AP</u>	RISING	<u>0.005MG/ML (0.005MG/ML)</u>	<u>A203897 002</u>	Nov 02, 2017
<u>AP</u>		<u>0.01MG/2ML (0.005MG/ML)</u>	<u>A203897 003</u>	Nov 02, 2017
<u>AP</u>	SANDOZ	<u>0.002MG/ML (0.002MG/ML)</u>	<u>A091108 001</u>	Jul 27, 2011
<u>AP</u>		<u>0.005MG/ML (0.005MG/ML)</u>	<u>A091108 002</u>	Jul 27, 2011
<u>AP</u>		<u>0.01MG/2ML (0.005MG/ML)</u>	<u>A091108 003</u>	Jul 27, 2011

ZEMPLAR

<u>AP</u>	+!	ABEVIE	<u>0.002MG/ML (0.002MG/ML)</u>	<u>N020819 002</u>	Feb 01, 2000
<u>AP</u>	+!		<u>0.005MG/ML (0.005MG/ML)</u>	<u>N020819 001</u>	Apr 17, 1998
<u>AP</u>	+!		<u>0.01MG/2ML (0.005MG/ML)</u>	<u>N020819 003</u>	Feb 01, 2000

PAROMOMYCIN SULFATE

CAPSULE; ORAL

PAROMOMYCIN SULFATE

! HERITAGE

EQ 250MG BASE

A065173 001 Dec 14, 2007

PAROXETINE HYDROCHLORIDE

SUSPENSION; ORAL

PAROXETINE HYDROCHLORIDE

! NOVITIUM PHARMA

EQ 10MG BASE/5ML

A215003 001 Sep 03, 2021

TABLET; ORAL

PAROXETINE

<u>AB</u>		PRINSTON INC	<u>EQ 10MG BASE</u>	<u>A203854 001</u>	Oct 31, 2014
<u>AB</u>			<u>EQ 20MG BASE</u>	<u>A203854 002</u>	Oct 31, 2014
<u>AB</u>			<u>EQ 30MG BASE</u>	<u>A203854 003</u>	Oct 31, 2014
<u>AB</u>			<u>EQ 40MG BASE</u>	<u>A203854 004</u>	Oct 31, 2014

PAROXETINE HYDROCHLORIDE

<u>AB</u>		APOTEX	<u>EQ 10MG BASE</u>	<u>A075356 001</u>	Jul 30, 2003
<u>AB</u>			<u>EQ 20MG BASE</u>	<u>A075356 002</u>	Jul 30, 2003
<u>AB</u>			<u>EQ 30MG BASE</u>	<u>A075356 003</u>	Jul 30, 2003
<u>AB</u>			<u>EQ 40MG BASE</u>	<u>A075356 004</u>	Jul 30, 2003
<u>AB</u>		AUROBINDO PHARMA	<u>EQ 10MG BASE</u>	<u>A078406 001</u>	Jul 25, 2007
<u>AB</u>			<u>EQ 20MG BASE</u>	<u>A078406 002</u>	Jul 25, 2007
<u>AB</u>			<u>EQ 30MG BASE</u>	<u>A078406 003</u>	Jul 25, 2007
<u>AB</u>			<u>EQ 40MG BASE</u>	<u>A078406 004</u>	Jul 25, 2007
<u>AB</u>		CHARTWELL RX	<u>EQ 10MG BASE</u>	<u>A076618 001</u>	Aug 15, 2005
<u>AB</u>			<u>EQ 20MG BASE</u>	<u>A076618 002</u>	Aug 15, 2005
<u>AB</u>			<u>EQ 30MG BASE</u>	<u>A076618 003</u>	Aug 15, 2005
<u>AB</u>			<u>EQ 40MG BASE</u>	<u>A076618 004</u>	Aug 15, 2005
<u>AB</u>		MYLAN	<u>EQ 10MG BASE</u>	<u>A078902 001</u>	Mar 13, 2008
<u>AB</u>			<u>EQ 20MG BASE</u>	<u>A078902 002</u>	Mar 13, 2008
<u>AB</u>			<u>EQ 30MG BASE</u>	<u>A078902 003</u>	Mar 13, 2008
<u>AB</u>			<u>EQ 40MG BASE</u>	<u>A078902 004</u>	Mar 13, 2008
<u>AB</u>		OXFORD PHARMS	<u>EQ 10MG BASE</u>	<u>A076968 001</u>	Jun 21, 2010
<u>AB</u>			<u>EQ 20MG BASE</u>	<u>A076968 002</u>	Jun 21, 2010
<u>AB</u>			<u>EQ 30MG BASE</u>	<u>A076968 003</u>	Jun 21, 2010
<u>AB</u>			<u>EQ 40MG BASE</u>	<u>A076968 004</u>	Jun 21, 2010
<u>AB</u>		YILING	<u>EQ 10MG BASE</u>	<u>A211248 001</u>	Nov 02, 2021
<u>AB</u>			<u>EQ 20MG BASE</u>	<u>A211248 002</u>	Nov 02, 2021
<u>AB</u>			<u>EQ 30MG BASE</u>	<u>A211248 003</u>	Nov 02, 2021
<u>AB</u>			<u>EQ 40MG BASE</u>	<u>A211248 004</u>	Nov 02, 2021
<u>AB</u>		ZYDUS PHARMS USA	<u>EQ 10MG BASE</u>	<u>A077584 001</u>	Mar 07, 2007
<u>AB</u>			<u>EQ 20MG BASE</u>	<u>A077584 002</u>	Mar 07, 2007
<u>AB</u>			<u>EQ 30MG BASE</u>	<u>A077584 003</u>	Mar 07, 2007
<u>AB</u>			<u>EQ 40MG BASE</u>	<u>A077584 004</u>	Mar 07, 2007

PAXIL

<u>AB</u>	+	APOTEX	<u>EQ 10MG BASE</u>	<u>N020031 001</u>	Dec 29, 1992
<u>AB</u>	+		<u>EQ 20MG BASE</u>	<u>N020031 002</u>	Dec 29, 1992
<u>AB</u>	+		<u>EQ 30MG BASE</u>	<u>N020031 003</u>	Dec 29, 1992
<u>AB</u>	+!		<u>EQ 40MG BASE</u>	<u>N020031 005</u>	Dec 29, 1992

TABLET, EXTENDED RELEASE; ORAL

PAROXETINE HYDROCHLORIDE

<u>AB</u>		AUROBINDO PHARMA USA	<u>EQ 12.5MG BASE</u>	<u>A077873 001</u>	Jun 29, 2007
<u>AB</u>			<u>EQ 25MG BASE</u>	<u>A077873 002</u>	Jun 29, 2007
<u>AB</u>			<u>EQ 37.5MG BASE</u>	<u>A091427 001</u>	Apr 14, 2011
<u>AB</u>		CADILA PHARMS LTD	<u>EQ 12.5MG BASE</u>	<u>A212645 001</u>	Aug 27, 2021
<u>AB</u>			<u>EQ 25MG BASE</u>	<u>A212645 002</u>	Aug 27, 2021
<u>AB</u>			<u>EQ 37.5MG BASE</u>	<u>A212645 003</u>	Aug 27, 2021
<u>AB</u>		CSPC OUYI	<u>EQ 12.5MG BASE</u>	<u>A213485 001</u>	Feb 16, 2021
<u>AB</u>			<u>EQ 25MG BASE</u>	<u>A213485 002</u>	Feb 16, 2021

PRESCRIPTION DRUG PRODUCT LIST

PAROXETINE HYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL

PAROXETINE HYDROCHLORIDE

<u>AB</u>		<u>EQ 37.5MG BASE</u>	<u>A213485 003</u>	Feb 16, 2021
<u>AB</u>	LANNETT CO INC	<u>EQ 12.5MG BASE</u>	<u>A204744 001</u>	Jun 10, 2016
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>A204744 002</u>	Jun 10, 2016
<u>AB</u>		<u>EQ 37.5MG BASE</u>	<u>A204744 003</u>	Jun 10, 2016
<u>AB</u>	LUPIN LTD	<u>EQ 12.5MG BASE</u>	<u>A204134 001</u>	Jan 20, 2017
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>A204134 002</u>	Jan 20, 2017
<u>AB</u>		<u>EQ 37.5MG BASE</u>	<u>A204134 003</u>	Jan 20, 2017
<u>AB</u>	PRINSTON INC	<u>EQ 12.5MG BASE</u>	<u>A209748 001</u>	Jan 04, 2024
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>A209748 002</u>	Jan 04, 2024
<u>AB</u>	SCIECURE PHARMA INC	<u>EQ 12.5MG BASE</u>	<u>A209293 001</u>	Jun 12, 2018
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>A209293 002</u>	Jun 12, 2018
<u>AB</u>		<u>EQ 37.5MG BASE</u>	<u>A209293 003</u>	Jun 12, 2018
	<u>PAXIL CR</u>			
<u>AB</u>	+ APOTEX	<u>EQ 12.5MG BASE</u>	<u>N020936 001</u>	Feb 16, 1999
<u>AB</u>	+	<u>EQ 25MG BASE</u>	<u>N020936 002</u>	Feb 16, 1999
<u>AB</u>	+!	<u>EQ 37.5MG BASE</u>	<u>N020936 003</u>	Dec 06, 2000

PAROXETINE MESYLATE

CAPSULE;ORAL

BRISDELLE

<u>AB</u>	+! LEGACY PHARMA	<u>EQ 7.5MG BASE</u>	<u>N204516 001</u>	Jun 28, 2013
-----------	------------------	----------------------	--------------------	--------------

PAROXETINE MESYLATE

<u>AB</u>	ACTAVIS LABS FL INC	<u>EQ 7.5MG BASE</u>	<u>A207139 001</u>	Jun 20, 2017
<u>AB</u>	PRINSTON INC	<u>EQ 7.5MG BASE</u>	<u>A207188 001</u>	Aug 18, 2017

PASIREOTIDE DIASPARTATE

SOLUTION;SUBCUTANEOUS

SIGNIFOR

+	RECORDATI RARE	EQ 0.3MG BASE/ML (EQ 0.3MG BASE/ML)	N200677 001	Dec 14, 2012
+		EQ 0.6MG BASE/ML (EQ 0.6MG BASE/ML)	N200677 002	Dec 14, 2012
+	!	EQ 0.9MG BASE/ML (EQ 0.9MG BASE/ML)	N200677 003	Dec 14, 2012

PASIREOTIDE PAMOATE

FOR SUSPENSION;INTRAMUSCULAR

SIGNIFOR LAR KIT

+	RECORDATI RARE	EQ 10MG BASE/VIAL	N203255 004	Jun 29, 2018
+		EQ 20MG BASE/VIAL	N203255 001	Dec 15, 2014
+		EQ 30MG BASE/VIAL	N203255 005	Jun 29, 2018
+		EQ 40MG BASE/VIAL	N203255 002	Dec 15, 2014
+	!	EQ 60MG BASE/VIAL	N203255 003	Dec 15, 2014

PATROMER SORBITEX CALCIUM

POWDER;ORAL

VELTASSA

+	VIFOR PHARMA	EQ 1GM BASE/PACKET	N205739 004	Oct 02, 2023
+		EQ 8.4GM BASE/PACKET	N205739 001	Oct 21, 2015
+	!	EQ 16.8GM BASE/PACKET	N205739 002	Oct 21, 2015

PATISIRAN SODIUM

SOLUTION;INTRAVENOUS

ONPATRO

+	ALNYLAM PHARMS INC	EQ 10MG BASE/5ML (EQ 2MG BASE/ML)	N210922 001	Aug 10, 2018
---	--------------------	-----------------------------------	-------------	--------------

PAZOPANIB HYDROCHLORIDE

TABLET;ORAL

PAZOPANIB HYDROCHLORIDE

<u>AB</u>	APOTEX	<u>EQ 200MG BASE</u>	<u>A217713 001</u>	Oct 19, 2023
<u>AB</u>	EUGIA PHARMA	<u>EQ 200MG BASE</u>	<u>A219034 001</u>	Dec 04, 2024
<u>AB</u>	NOVUGEN	<u>EQ 200MG BASE</u>	<u>A218231 001</u>	Apr 23, 2024
<u>AB</u>	SUN PHARM	<u>EQ 200MG BASE</u>	<u>A215837 001</u>	Oct 19, 2023
<u>AB</u>	TEVA PHARMS INC	<u>EQ 200MG BASE</u>	<u>A217517 001</u>	Oct 19, 2023
	<u>VOTRIENT</u>			
<u>AB</u>	+! NOVARTIS	<u>EQ 200MG BASE</u>	<u>N022465 001</u>	Oct 19, 2009

PEGCETACOPLAN

SOLUTION;INTRAVITREAL

SYFOVRE

+	APELLIS PHARMS	15MG/0.1ML (15MG/0.1ML)	N217171 001	Feb 17, 2023
---	----------------	-------------------------	-------------	--------------

SOLUTION;SUBCUTANEOUS

EMPAVELI

+	APELLIS PHARMS	1080MG/20ML (54MG/ML)	N215014 001	May 14, 2021
---	----------------	-----------------------	-------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

PEGULICIANINE ACETATE

POWDER; INTRAVENOUS

LUMISIGHT

+! LUMICELL EQ 40MG BASE/VIAL N214511 001 Apr 17, 2024

PEMETREXED

SOLUTION; INTRAVENOUS

PEMETREXED

+! ACTAVIS 100MG/4ML (25MG/ML) N208419 001 Aug 21, 2020

+! 500MG/20ML (25MG/ML) N208419 002 Aug 21, 2020

+! 1GM/40ML (25MG/ML) N208419 003 Aug 21, 2020

PEMFEXY

+! EAGLE PHARMS 500MG/20ML (25MG/ML) N209472 001 Feb 08, 2020

PEMETREXED DIPOTASSIUM

POWDER; INTRAVENOUS

AXTLE

+! AVYXA HOLDINGS EQ 100MG BASE/VIAL N210661 001 Jun 28, 2024

+! EQ 500MG BASE/VIAL N210661 002 Jun 28, 2024

PEMETREXED DISODIUM

POWDER; INTRAVENOUS

ALIMTAAP +! LILLY EQ 100MG BASE/VIAL N021462 002 Sep 07, 2007AP +! EQ 500MG BASE/VIAL N021462 001 Feb 04, 2004PEMETREXED DISODIUMAP ACCORD HLTHCARE EQ 100MG BASE/VIAL A203485 001 May 25, 2022AP EQ 500MG BASE/VIAL A203485 002 May 25, 2022AP EQ 1GM BASE/VIAL A203485 003 May 25, 2022AP APOTEX EQ 100MG BASE/VIAL A203774 001 May 25, 2022AP EQ 500MG BASE/VIAL A203774 002 May 25, 2022AP EQ 750MG BASE/VIAL A209851 001 May 25, 2022AP EQ 1GM BASE/VIAL A209085 001 May 25, 2022AP BAXTER HLTHCARE EQ 100MG BASE/VIAL A214436 001 Aug 18, 2022

CORP

AP EQ 500MG BASE/VIAL A214436 002 Aug 18, 2022AP DR REDDYS EQ 100MG BASE/VIAL A202596 001 May 25, 2022AP EQ 500MG BASE/VIAL A202596 002 May 25, 2022AP EQ 1GM BASE/VIAL A202596 003 May 25, 2022AP EUGIA PHARMA EQ 100MG BASE/VIAL A214632 001 May 25, 2022AP EQ 500MG BASE/VIAL A214632 002 May 25, 2022AP EQ 1GM BASE/VIAL A214632 003 May 25, 2022AP FRESENIUS KABI USA EQ 100MG BASE/VIAL A090384 001 May 25, 2022AP EQ 500MG BASE/VIAL A090384 002 May 25, 2022AP ! EQ 750MG BASE/VIAL A090384 003 May 25, 2022AP ! EQ 1GM BASE/VIAL A090384 004 May 25, 2022AP HETERO LABS LTD VI EQ 100MG BASE/VIAL A215460 001 Jun 14, 2024AP EQ 500MG BASE/VIAL A215460 002 Jun 14, 2024AP JIANGSU HANSO EQ 100MG BASE/VIAL A208696 001 May 25, 2022

PHARM

AP EQ 500MG BASE/VIAL A208696 002 May 25, 2022AP MEITHEAL EQ 100MG BASE/VIAL A215479 001 Dec 13, 2022AP EQ 500MG BASE/VIAL A215479 002 Dec 13, 2022AP EQ 750MG BASE/VIAL A215479 003 Dec 13, 2022AP EQ 1GM BASE/VIAL A215479 004 Dec 13, 2022AP NANG KUANG PHARM CO EQ 100MG BASE/VIAL A207352 001 May 25, 2022AP EQ 500MG BASE/VIAL A207352 002 May 25, 2022AP PRINSTON INC EQ 100MG BASE/VIAL A216582 001 Dec 11, 2023AP QILU PHARM HAINAN EQ 100MG BASE/VIAL A204890 001 May 25, 2022AP EQ 500MG BASE/VIAL A204890 002 May 25, 2022AP RELIANCE LIFE EQ 100MG BASE/VIAL A211899 001 May 25, 2022AP EQ 500MG BASE/VIAL A211899 002 May 25, 2022AP ZYDUS PHARMS EQ 100MG BASE/VIAL A214073 001 May 25, 2022AP EQ 500MG BASE/VIAL A214073 002 May 25, 2022AP EQ 1GM BASE/VIAL A214073 003 May 25, 2022

SOLUTION; INTRAVENOUS

PEMETREXED

+! SHILPA EQ 100MG/10ML BASE (10MG/ML) N215179 001 May 22, 2023

+! EQ 500MG/50ML BASE (10MG/ML) N215179 002 May 22, 2023

+! EQ 1GM/100ML BASE (10MG/ML) N215179 003 May 22, 2023

PEMETREXED DISODIUM

+! ACCORD HLTHCARE EQ 100MG BASE/4ML (EQ 25MG BASE/ML) N214408 001 Jul 19, 2022

+! EQ 500MG BASE/20ML (EQ 25MG BASE/ML) N214408 002 Jul 19, 2022

+! EQ 850MG BASE/34ML (EQ 25MG BASE/ML) N214408 003 Jul 19, 2022

PRESCRIPTION DRUG PRODUCT LIST

PEMETREXED DISODIUM

SOLUTION; INTRAVENOUS

PEMETREXED DISODIUM

+	!		EQ 1GM BASE/40ML (EQ 25MG BASE/ML)	N214408	004	Jul 19, 2022
+	!	HOSPIRA	EQ 100MG BASE/4ML (EQ 25MG BASE/ML)	N214218	001	Jun 22, 2022
+	!		EQ 500MG BASE/20ML (EQ 25MG BASE/ML)	N214218	002	Jun 22, 2022
+	!		EQ 1GM BASE/40ML (EQ 25MG BASE/ML)	N214218	003	Jun 22, 2022
+	!	SANDOZ	EQ 100MG BASE/4ML (EQ 25MG BASE/ML)	N214657	001	May 26, 2022
+	!		EQ 500MG BASE/20ML (EQ 25MG BASE/ML)	N214657	002	May 26, 2022
+	!		EQ 1GM BASE/40ML (EQ 25MG BASE/ML)	N214657	003	May 26, 2022

PEMETREXED DITROMETHAMINE

POWDER; INTRAVENOUS

PEMETREXED DITROMETHAMINE

+	!	HOSPIRA	EQ 100MG BASE/VIAL	N208746	001	Jun 10, 2022
+	!		EQ 500MG BASE/VIAL	N208746	002	Jun 10, 2022

PEMIGATINIB

TABLET; ORAL

PEMAZYRE

+		INCYTE CORP	4.5MG	N213736	001	Apr 17, 2020
+			9MG	N213736	002	Apr 17, 2020
+	!		13.5MG	N213736	003	Apr 17, 2020

PENCICLOVIR

CREAM; TOPICAL

DENAVIR

AB	+	!	MYLAN	1%	N020629	001	Sep 24, 1996
-----------	---	---	-------	-----------	----------------	------------	--------------

PENCICLOVIR

AB			PADAGIS ISRAEL	1%	A212368	001	Sep 27, 2024
AB			TEVA PHARMS USA	1%	A212710	001	Nov 09, 2022
AB			TORRENT	1%	A216981	001	Aug 07, 2023

PENICILLAMINE

CAPSULE; ORAL

CUPRIMINE

AB	+	!	VALEANT PHARMS INTL	250MG	N019853	001	
-----------	---	---	---------------------	--------------	----------------	------------	--

PENICILLAMINE

AB			APOTEX	250MG	A213310	001	Apr 28, 2020
AB			BRECKENRIDGE	250MG	A215409	001	Aug 23, 2021
AB			DR REDDYS	250MG	A211867	001	Aug 04, 2020
AB			ENDO OPERATIONS	250MG	A211231	001	Dec 23, 2019
AB			GRANULES	250MG	A211735	001	Dec 02, 2020
AB			INVAGEN PHARMS	250MG	A213293	001	Aug 19, 2021
AB			NAVINTA LLC	250MG	A214363	001	Oct 08, 2021
AB			WATSON LABS INC	250MG	A210976	001	Jun 24, 2019

TABLET; ORAL

DEPEN

AB	+	!	MYLAN SPECIALITY LP	250MG	N019854	001	
-----------	---	---	---------------------	--------------	----------------	------------	--

PENICILLAMINE

AB			ENDO OPERATIONS	250MG	A211196	001	Dec 23, 2019
AB			LUPIN LTD	250MG	A212933	001	Nov 30, 2020

PENICILLIN G BENZATHINE

INJECTABLE; INJECTION

BICILLIN L-A

BC	+	!	KING PHARMS LLC	600,000 UNITS/ML	N050141	001	
----	---	---	-----------------	------------------	---------	-----	--

PENICILLIN G BENZATHINE; PENICILLIN G PROCAINE

INJECTABLE; INJECTION

BICILLIN C-R

+	!		KING PHARMS LLC	300,000 UNITS/ML; 300,000 UNITS/ML	N050138	001	
---	---	--	-----------------	------------------------------------	---------	-----	--

BICILLIN C-R 900/300

+	!		KING PHARMS LLC	900,000 UNITS/2ML; 300,000 UNITS/2ML	N050138	003	
---	---	--	-----------------	--------------------------------------	---------	-----	--

PENICILLIN G POTASSIUM

INJECTABLE; INJECTION

PENICILLIN G POTASSIUM

AP			ACS DOBFAR SPA	20,000,000 UNITS/VIAL	A205043	002	Oct 26, 2018
AP				5,000,000 UNITS/VIAL	A205043	001	Oct 26, 2018
AP			HQ SPECLT PHARMA	5,000,000 UNITS/VIAL	A065149	002	Jul 23, 2009
AP				20,000,000 UNITS/VIAL	A065149	003	Jul 23, 2009
AP			ISTITUTO BIO ITA SPA	5,000,000 UNITS/VIAL	A065448	001	Aug 18, 2009
AP				20,000,000 UNITS/VIAL	A065448	002	Aug 18, 2009
AP			SANDOZ	5,000,000 UNITS/VIAL	A065079	002	Aug 30, 2002

PRESCRIPTION DRUG PRODUCT LIST

PENICILLIN G POTASSIUM

INJECTABLE; INJECTION

PENICILLIN G POTASSIUM

AP		<u>20,000,000 UNITS/VIAL</u>	<u>A065079 003</u>	Aug 30, 2002
-----------	--	-------------------------------------	---------------------------	--------------

PFIZERPEN

AP	!	PFIZER	<u>5,000,000 UNITS/VIAL</u>	<u>A060657 002</u>	
-----------	---	--------	------------------------------------	---------------------------	--

AP	!		<u>20,000,000 UNITS/VIAL</u>	<u>A060657 003</u>	
-----------	---	--	-------------------------------------	---------------------------	--

PENICILLIN G POTASSIUM

HQ SPECLT PHARMA 1,000,000 UNITS/VIAL

A065149 001 Jul 23, 2009

PENICILLIN G POTASSIUM IN PLASTIC CONTAINER

+! BAXTER HLTHCARE 20,000 UNITS/ML

N050638 001 Jun 25, 1990

+! 40,000 UNITS/ML

N050638 002 Jun 25, 1990

+! 60,000 UNITS/ML

N050638 003 Jun 25, 1990

PENICILLIN G PROCAINE

INJECTABLE; INJECTION

PENICILLIN G PROCAINE

! KING PHARMS LLC 300,000 UNITS/ML

A060101 002

! 600,000 UNITS/ML

A060101 001

PENICILLIN G SODIUM

INJECTABLE; INTRAMUSCULAR, INTRAVENOUS

PENICILLIN G SODIUM

! SANDOZ 5,000,000 UNITS/VIAL

A065068 001 Feb 26, 2001

PENICILLIN V POTASSIUM

FOR SOLUTION; ORAL

PENICILLIN-VK

TEVA EQ 125MG BASE/5ML

A060456 001

! EQ 250MG BASE/5ML

A060456 002

TABLET; ORAL

PENICILLIN V POTASSIUM

AB		AUROBINDO PHARMA	<u>EQ 250MG BASE</u>	<u>A065435 001</u>	Apr 29, 2008
-----------	--	------------------	-----------------------------	---------------------------	--------------

AB	!		<u>EQ 500MG BASE</u>	<u>A065435 002</u>	Apr 29, 2008
-----------	---	--	-----------------------------	---------------------------	--------------

AB		CHARTWELL RX	<u>EQ 250MG BASE</u>	<u>A062936 001</u>	Nov 25, 1988
-----------	--	--------------	-----------------------------	---------------------------	--------------

AB			<u>EQ 500MG BASE</u>	<u>A062935 001</u>	Nov 23, 1988
-----------	--	--	-----------------------------	---------------------------	--------------

AB		HIKMA PHARMS	<u>EQ 250MG BASE</u>	<u>A090549 001</u>	Oct 11, 2013
-----------	--	--------------	-----------------------------	---------------------------	--------------

AB			<u>EQ 500MG BASE</u>	<u>A090549 002</u>	Oct 11, 2013
-----------	--	--	-----------------------------	---------------------------	--------------

PENICILLIN-VK

AB		TEVA	<u>EQ 250MG BASE</u>	<u>A060711 002</u>	
-----------	--	------	-----------------------------	---------------------------	--

AB			<u>EQ 500MG BASE</u>	<u>A060711 003</u>	
-----------	--	--	-----------------------------	---------------------------	--

PENTAMIDINE ISETHIONATE

FOR SOLUTION; INHALATION

NEBUPENT

AN	+!	FRESENIUS KABI USA	<u>300MG/VIAL</u>	<u>N019887 001</u>	Jun 15, 1989
-----------	----	--------------------	--------------------------	---------------------------	--------------

PENTAMIDINE ISETHIONATE

AN		SETON PHARMS	<u>300MG/VIAL</u>	<u>A206667 001</u>	Apr 24, 2019
-----------	--	--------------	--------------------------	---------------------------	--------------

AN		X-GEN PHARMS INC	<u>300MG/VIAL</u>	<u>A206983 001</u>	Jan 20, 2023
-----------	--	------------------	--------------------------	---------------------------	--------------

INJECTABLE; INJECTION

PENTAM

AP	+!	FRESENIUS KABI USA	<u>300MG/VIAL</u>	<u>N019264 001</u>	Oct 16, 1984
-----------	----	--------------------	--------------------------	---------------------------	--------------

PENTAMIDINE ISETHIONATE

AP		AVET LIFESCIENCES	<u>300MG/VIAL</u>	<u>A213806 001</u>	Jan 07, 2021
-----------	--	-------------------	--------------------------	---------------------------	--------------

AP		SETON PHARMS	<u>300MG/VIAL</u>	<u>A206666 001</u>	Sep 28, 2017
-----------	--	--------------	--------------------------	---------------------------	--------------

AP		XGEN PHARMS	<u>300MG/VIAL</u>	<u>A206982 001</u>	Mar 17, 2022
-----------	--	-------------	--------------------------	---------------------------	--------------

PENTOBARBITAL SODIUM

INJECTABLE; INJECTION

NEMBUTAL SODIUM

AP	+!	RISING	<u>50MG/ML</u>	<u>A083246 001</u>	
-----------	----	--------	-----------------------	---------------------------	--

PENTOBARBITAL SODIUM

AP		BPI LABS	<u>50MG/ML</u>	<u>A206677 001</u>	Nov 27, 2017
-----------	--	----------	-----------------------	---------------------------	--------------

AP		HIKMA	<u>50MG/ML</u>	<u>A203619 001</u>	Nov 13, 2017
-----------	--	-------	-----------------------	---------------------------	--------------

AP		SAGENT PHARMS INC	<u>50MG/ML</u>	<u>A206404 001</u>	May 23, 2016
-----------	--	-------------------	-----------------------	---------------------------	--------------

PENTOSAN POLYSULFATE SODIUM

CAPSULE; ORAL

ELMIRON

+! JANSSEN PHARMS 100MG

N020193 001 Sep 26, 1996

PRESCRIPTION DRUG PRODUCT LIST

PENTOSTATIN

INJECTABLE; INJECTION

NIPENT

AP	+ !	HOSPIRA INC	10MG/VIAL	N020122 001	Oct 11, 1991
-----------	------------	-------------	------------------	--------------------	--------------

PENTOSTATIN

AP		WEST-WARD PHARMS INT	10MG/VIAL	A077841 001	Aug 07, 2007
-----------	--	-------------------------	------------------	--------------------	--------------

PENTOXIFYLLINE

TABLET, EXTENDED RELEASE; ORAL

PENTOXIFYLLINE

AB		ANI PHARMS	400MG	A074878 001	Jul 09, 1997
AB	!	APOTEX	400MG	A075191 001	Jun 09, 1999
AB		RISING	400MG	A074425 001	Jul 08, 1997
AB		VALEANT PHARMS	400MG	A075028 001	Jul 20, 1998

PERAMIVIR

SOLUTION; INTRAVENOUS

RAPIVAB

+ !	BIOCRYS	200MG/20ML (10MG/ML)	N206426 001	Dec 19, 2014
------------	---------	----------------------	-------------	--------------

PERAMPANEL

SUSPENSION; ORAL

FYCOMPA

+ !	CATALYST PHARMS	0.5MG/ML	N208277 001	Apr 29, 2016
------------	-----------------	----------	-------------	--------------

TABLET; ORAL

FYCOMPA

+	CATALYST PHARMS	2MG	N202834 001	Oct 22, 2012
+		4MG	N202834 002	Oct 22, 2012
+		6MG	N202834 003	Oct 22, 2012
+		8MG	N202834 004	Oct 22, 2012
+		10MG	N202834 005	Oct 22, 2012
+ !		12MG	N202834 006	Oct 22, 2012

PERFLUOROHEXYLOCTANE

SOLUTION/DROPS; OPHTHALMIC

MIEBO

+ !	BAUSCH AND LOMB INC	1.338GM/ML	N216675 001	May 18, 2023
------------	---------------------	------------	-------------	--------------

PERFLUTREN

INJECTABLE; INTRAVENOUS

DEFINITY

+ !	LANTHEUS MEDCL	13.04MG/2ML (6.52MG/ML)	N021064 001	Jul 31, 2001
------------	----------------	-------------------------	-------------	--------------

DEFINITY RT

+ !	LANTHEUS MEDCL	13.04MG/2ML (6.52MG/ML)	N021064 002	Nov 17, 2020
------------	----------------	-------------------------	-------------	--------------

PERINDOPRIL ERBUMINE

TABLET; ORAL

PERINDOPRIL ERBUMINE

	AUROBINDO PHARMA	2MG	A079070 001	Nov 10, 2009
--	------------------	-----	-------------	--------------

		4MG	A079070 002	Nov 10, 2009
--	--	-----	-------------	--------------

!		8MG	A079070 003	Nov 10, 2009
----------	--	-----	-------------	--------------

PERMETHRIN

CREAM; TOPICAL

ELIMITE

AB	+	AUROBINDO PHARMA USA	5%	N019855 001	Aug 25, 1989
-----------	----------	-------------------------	-----------	--------------------	--------------

PERMETHRIN

AB		ACTAVIS LABS	5%	A074806 001	Jan 23, 1998
AB		ENCUBE ETHICALS	5%	A211303 001	Apr 03, 2019
AB	!	PADAGIS ISRAEL	5%	A076369 001	Apr 21, 2003

PERPHENAZINE

TABLET; ORAL

PERPHENAZINE

AB		APPCO	2MG	A210163 001	May 18, 2022
AB			4MG	A210163 002	May 18, 2022
AB			8MG	A210163 003	May 18, 2022
AB			16MG	A210163 004	May 18, 2022
AB		ENDO OPERATIONS	2MG	A040226 001	Dec 31, 1998
AB			4MG	A040226 002	Dec 31, 1998
AB			8MG	A040226 003	Dec 31, 1998
AB			16MG	A040226 004	Dec 31, 1998
AB		MACLEODS PHARMS LTD	2MG	A212545 001	May 06, 2024
AB			4MG	A212545 002	May 06, 2024
AB			8MG	A212545 003	May 06, 2024

PRESCRIPTION DRUG PRODUCT LIST

PERPHENAZINE

TABLET; ORAL

PERPHENAZINE

<u>AB</u>		<u>16MG</u>	<u>A212545</u>	<u>004</u>	May 06, 2024
<u>AB</u>	RISING	<u>2MG</u>	<u>A205056</u>	<u>001</u>	Mar 01, 2019
<u>AB</u>		<u>4MG</u>	<u>A205056</u>	<u>002</u>	Mar 01, 2019
<u>AB</u>		<u>8MG</u>	<u>A205056</u>	<u>003</u>	Mar 01, 2019
<u>AB</u>		<u>16MG</u>	<u>A205056</u>	<u>004</u>	Mar 01, 2019
<u>AB</u>	SANDOZ	<u>2MG</u>	<u>A089685</u>	<u>002</u>	Dec 08, 1988
<u>AB</u>		<u>4MG</u>	<u>A089685</u>	<u>003</u>	Dec 08, 1988
<u>AB</u>		<u>8MG</u>	<u>A089685</u>	<u>001</u>	Dec 08, 1988
<u>AB</u>	!	<u>16MG</u>	<u>A089685</u>	<u>004</u>	Dec 08, 1988
<u>AB</u>	WATSON LABS INC	<u>2MG</u>	<u>A207582</u>	<u>001</u>	Oct 17, 2016
<u>AB</u>		<u>4MG</u>	<u>A207582</u>	<u>002</u>	Oct 17, 2016
<u>AB</u>		<u>8MG</u>	<u>A207582</u>	<u>003</u>	Oct 17, 2016
<u>AB</u>		<u>16MG</u>	<u>A207582</u>	<u>004</u>	Oct 17, 2016
<u>AB</u>	WILSHIRE PHARMS INC	<u>2MG</u>	<u>A205973</u>	<u>001</u>	Dec 17, 2015
<u>AB</u>		<u>4MG</u>	<u>A205973</u>	<u>002</u>	Dec 17, 2015
<u>AB</u>		<u>8MG</u>	<u>A205973</u>	<u>003</u>	Dec 17, 2015
<u>AB</u>		<u>16MG</u>	<u>A205973</u>	<u>004</u>	Dec 17, 2015
<u>AB</u>	ZYDUS PHARMS	<u>2MG</u>	<u>A205232</u>	<u>001</u>	Apr 06, 2020
<u>AB</u>		<u>4MG</u>	<u>A205232</u>	<u>002</u>	Apr 06, 2020
<u>AB</u>		<u>8MG</u>	<u>A205232</u>	<u>003</u>	Apr 06, 2020
<u>AB</u>		<u>16MG</u>	<u>A205232</u>	<u>004</u>	Apr 06, 2020

PEXIDARTINIB HYDROCHLORIDE

CAPSULE; ORAL

TURALIO

+ DAIICHI SANKYO INC EQ 125MG BASE N211810 002 Oct 14, 2022

PHENDIMETRAZINE TARTRATE

CAPSULE, EXTENDED RELEASE; ORAL

PHENDIMETRAZINE TARTRATE

+! VIRTUS 105MG N018074 001

TABLET; ORAL

BONTRIL PDMAA +! VALEANT 35MG A085272 001PHENDIMETRAZINE TARTRATE

<u>AA</u>	CHARTWELL	<u>35MG</u>	<u>A089452</u>	<u>001</u>	Oct 30, 1991
<u>AA</u>	ELITE LABS INC	<u>35MG</u>	<u>A040762</u>	<u>001</u>	Jan 28, 2008
<u>AA</u>	KVK TECH	<u>35MG</u>	<u>A091042</u>	<u>001</u>	Aug 31, 2010
<u>AA</u>	VIRTUS	<u>35MG</u>	<u>A085588</u>	<u>001</u>	

PHENELZINE SULFATE

TABLET; ORAL

NARDILAB +! PARKE DAVIS EQ 15MG BASE N011909 002PHENELZINE SULFATEAB NOVEL LABS INC EQ 15MG BASE A200181 001 Dec 08, 2010PHENOBARBITAL SODIUM

POWDER; INTRAVENOUS

SEZABY

+! SUN PHARM INDS INC 100MG/VIAL N215910 001 Nov 17, 2022

PHENOXYBENZAMINE HYDROCHLORIDE

CAPSULE; ORAL

DIBENZYLINEAB + ADVANZ PHARMA 10MG N008708 001PHENOXYBENZAMINE HYDROCHLORIDE

<u>AB</u>	AMNEAL	<u>10MG</u>	<u>A212568</u>	<u>001</u>	Oct 27, 2020
<u>AB</u>	AUROBINDO PHARMA	<u>10MG</u>	<u>A215600</u>	<u>001</u>	May 08, 2023
<u>AB</u>	!	<u>10MG</u>	<u>A204522</u>	<u>001</u>	Jan 24, 2017
<u>AB</u>	NOVITIUM PHARMA	<u>10MG</u>	<u>A215042</u>	<u>001</u>	Jul 19, 2022
<u>AB</u>	RISING	<u>10MG</u>	<u>A204551</u>	<u>001</u>	Jun 20, 2023

PHENTERMINE HYDROCHLORIDE

CAPSULE; ORAL

PHENTERMINE HYDROCHLORIDE

<u>AA</u>	AUROLIFE PHARMA LLC	<u>15MG</u>	<u>A204318</u>	<u>001</u>	Nov 09, 2016
<u>AA</u>		<u>30MG</u>	<u>A204318</u>	<u>002</u>	Nov 09, 2016
<u>AA</u>	ELITE LABS	<u>15MG</u>	<u>A202248</u>	<u>001</u>	Sep 28, 2012
<u>AA</u>		<u>30MG</u>	<u>A202248</u>	<u>002</u>	Sep 28, 2012
<u>AA</u>	KVK TECH	<u>15MG</u>	<u>A040886</u>	<u>002</u>	Mar 31, 2008
<u>AA</u>		<u>30MG</u>	<u>A040875</u>	<u>001</u>	Mar 21, 2008

PRESCRIPTION DRUG PRODUCT LIST

PHENTERMINE HYDROCHLORIDE

CAPSULE; ORAL

PHENTERMINE HYDROCHLORIDE

<u>AA</u>		<u>30MG</u>	<u>A040886</u>	<u>001</u>	Mar 31, 2008
<u>AA</u>		<u>37.5MG</u>	<u>A040887</u>	<u>001</u>	Apr 24, 2008
<u>AA</u>	NUVO PHARM	<u>15MG</u>	<u>A205019</u>	<u>001</u>	Dec 05, 2014
<u>AA</u>		<u>30MG</u>	<u>A205019</u>	<u>002</u>	Dec 05, 2014
<u>AA</u>	!	<u>37.5MG</u>	<u>A205017</u>	<u>001</u>	Sep 25, 2014
<u>AA</u>	+!	<u>15MG</u>	<u>A087190</u>	<u>002</u>	
<u>AA</u>	+!	<u>30MG</u>	<u>A086945</u>	<u>001</u>	Jul 20, 1983
<u>AA</u>	+!	<u>30MG</u>	<u>A087190</u>	<u>001</u>	

TABLET; ORAL

ADIPEX-P

<u>AA</u>	+!	TEVA	<u>37.5MG</u>	<u>A085128</u>	<u>001</u>	
-----------	----	------	---------------	----------------	------------	--

LOMAIRA

<u>AA</u>	!	AVANTHI INC	<u>8MG</u>	<u>A203495</u>	<u>001</u>	Sep 12, 2016
-----------	---	-------------	------------	----------------	------------	--------------

PHENTERMINE HYDROCHLORIDE

<u>AA</u>		AUROBINDO PHARMA LTD	<u>37.5MG</u>	<u>A203068</u>	<u>001</u>	Aug 06, 2014
<u>AA</u>		ELITE LABS	<u>37.5MG</u>	<u>A200272</u>	<u>001</u>	Jan 31, 2011
<u>AA</u>		ELITE LABS INC	<u>37.5MG</u>	<u>A040190</u>	<u>001</u>	May 30, 1997
<u>AA</u>		KVK TECH	<u>37.5MG</u>	<u>A040876</u>	<u>001</u>	Mar 31, 2008
<u>AA</u>		KVK TECH INC	<u>8MG</u>	<u>A203436</u>	<u>001</u>	Mar 17, 2017
<u>AA</u>		MERRO PHARM USA	<u>37.5MG</u>	<u>A206342</u>	<u>001</u>	Nov 18, 2016
<u>AA</u>		NUVO PHARM	<u>37.5MG</u>	<u>A205008</u>	<u>001</u>	Sep 25, 2014
<u>AA</u>		PRINSTON INC	<u>37.5MG</u>	<u>A040377</u>	<u>001</u>	Jan 04, 2002
<u>AA</u>		SUN PHARM INDUSTRIES	<u>37.5MG</u>	<u>A040526</u>	<u>001</u>	Oct 23, 2003

TABLET, ORALLY DISINTEGRATING; ORAL

PHENTERMINE HYDROCHLORIDE

		ZYDUS PHARMS	15MG	A204663	001	Jun 28, 2017
			30MG	A204663	002	Jun 28, 2017
			37.5MG	A204663	003	Jun 28, 2017

PHENTERMINE HYDROCHLORIDE; TOPIRAMATE

CAPSULE, EXTENDED RELEASE; ORAL

PHENTERMINE HYDROCHLORIDE AND TOPIRAMATE

<u>AB</u>		ACTAVIS LABS FL INC	<u>EQ 3.75MG BASE;23MG</u>	<u>A204982</u>	<u>001</u>	Jun 25, 2024
<u>AB</u>			<u>EQ 7.5MG BASE;46MG</u>	<u>A204982</u>	<u>002</u>	Jun 25, 2024
<u>AB</u>			<u>EQ 11.25MG BASE;69MG</u>	<u>A204982</u>	<u>003</u>	Jun 25, 2024
<u>AB</u>			<u>EQ 15MG BASE;92MG</u>	<u>A204982</u>	<u>004</u>	Jun 25, 2024

OSYMIA

<u>AB</u>	+	VIVUS LLC	<u>EQ 3.75MG BASE;23MG</u>	<u>N022580</u>	<u>001</u>	Jul 17, 2012
<u>AB</u>	+		<u>EQ 7.5MG BASE;46MG</u>	<u>N022580</u>	<u>002</u>	Jul 17, 2012
<u>AB</u>	+		<u>EQ 11.25MG BASE;69MG</u>	<u>N022580</u>	<u>003</u>	Jul 17, 2012
<u>AB</u>	+!		<u>EQ 15MG BASE;92MG</u>	<u>N022580</u>	<u>004</u>	Jul 17, 2012

PHENTOLAMINE MESYLATE

INJECTABLE; INJECTION

PHENTOLAMINE MESYLATE

<u>AP</u>	!	HIKMA	<u>5MG/VIAL</u>	<u>A040235</u>	<u>001</u>	Mar 11, 1998
<u>AP</u>		PRECISION DOSE INC	<u>5MG/VIAL</u>	<u>A207686</u>	<u>001</u>	Jul 14, 2017

ORAVVERSE

	+!	SEPTODONT HOLDING	0.4MG/1.7ML	N022159	001	May 09, 2008
--	----	-------------------	-------------	---------	-----	--------------

SOLUTION; OPHTHALMIC

RYZUMVI

	+!	FAMYGEN LIFE SCI	EQ 0.75% BASE	N217064	001	Sep 25, 2023
--	----	------------------	---------------	---------	-----	--------------

PHENYLEPHRINE HYDROCHLORIDE

SOLUTION; INTRAVENOUS

PHENYLEPHRINE HYDROCHLORIDE

<u>AP1</u>		AMNEAL	<u>10MG/ML (10MG/ML)</u>	<u>A211079</u>	<u>001</u>	Jul 05, 2018
<u>AP1</u>			<u>50MG/5ML (10MG/ML)</u>	<u>A211078</u>	<u>001</u>	Jul 19, 2018
<u>AP1</u>			<u>100MG/10ML (10MG/ML)</u>	<u>A211078</u>	<u>002</u>	Jul 19, 2018
<u>AP1</u>		ASPIRO	<u>10MG/ML (10MG/ML)</u>	<u>A218110</u>	<u>001</u>	May 10, 2024
<u>AP1</u>			<u>50MG/5ML (10MG/ML)</u>	<u>A218110</u>	<u>002</u>	May 10, 2024
<u>AP1</u>			<u>100MG/10ML (10MG/ML)</u>	<u>A218110</u>	<u>003</u>	May 10, 2024
<u>AP1</u>		CAPLIN	<u>10MG/ML (10MG/ML)</u>	<u>A213318</u>	<u>001</u>	Jun 11, 2020
<u>AP1</u>			<u>50MG/5ML (10MG/ML)</u>	<u>A213318</u>	<u>002</u>	Jun 11, 2020
<u>AP1</u>			<u>100MG/10ML (10MG/ML)</u>	<u>A213318</u>	<u>003</u>	Jun 11, 2020
<u>AP1</u>		ENDO OPERATIONS	<u>50MG/5ML (10MG/ML)</u>	<u>A210025</u>	<u>002</u>	Dec 21, 2018
<u>AP1</u>			<u>100MG/10ML (10MG/ML)</u>	<u>A210025</u>	<u>003</u>	Dec 21, 2018
<u>AP1</u>		EUGIA PHARMA	<u>50MG/5ML (10MG/ML)</u>	<u>A210697</u>	<u>001</u>	Nov 13, 2020
<u>AP1</u>			<u>100MG/10ML (10MG/ML)</u>	<u>A210697</u>	<u>002</u>	Nov 13, 2020

PRESCRIPTION DRUG PRODUCT LIST

PHENYLEPHRINE HYDROCHLORIDE

SOLUTION; INTRAVENOUS

PHENYLEPHRINE HYDROCHLORIDE

<u>AP1</u>	FRESENIUS KABI USA	<u>50MG/5ML (10MG/ML)</u>	<u>A210666 001</u>	Jan 30, 2019
<u>AP1</u>		<u>100MG/10ML (10MG/ML)</u>	<u>A210666 002</u>	Jan 30, 2019
<u>AP1</u>	GLAND PHARMA LTD	<u>10MG/ML (10MG/ML)</u>	<u>A211920 003</u>	Apr 08, 2021
<u>AP1</u>		<u>50MG/5ML (10MG/ML)</u>	<u>A211920 001</u>	Jun 05, 2020
<u>AP1</u>		<u>100MG/10ML (10MG/ML)</u>	<u>A211920 002</u>	Jun 05, 2020
<u>AP1</u>	MANKIND PHARMA	<u>10MG/ML (10MG/ML)</u>	<u>A217069 001</u>	Aug 30, 2022
<u>AP1</u>		<u>50MG/5ML (10MG/ML)</u>	<u>A217069 002</u>	Aug 30, 2022
<u>AP1</u>		<u>100MG/10ML (10MG/ML)</u>	<u>A217069 003</u>	Aug 30, 2022
<u>AP1</u>	MEITHEAL	<u>50MG/5ML (10MG/ML)</u>	<u>A210333 001</u>	Apr 27, 2018
<u>AP1</u>		<u>100MG/10ML (10MG/ML)</u>	<u>A210333 002</u>	Apr 27, 2018
<u>AP1</u>	PROVEPHARM SAS	<u>10MG/ML (10MG/ML)</u>	<u>A211081 001</u>	Jul 17, 2020
<u>AP1</u>		<u>50MG/5ML (10MG/ML)</u>	<u>A211081 002</u>	Jul 17, 2020
<u>AP1</u>		<u>100MG/10ML (10MG/ML)</u>	<u>A211081 003</u>	Jul 17, 2020
<u>AP1</u>	SAGENT PHARMS INC	<u>10MG/ML (10MG/ML)</u>	<u>A209967 001</u>	Jan 16, 2020
<u>AP1</u>		<u>50MG/5ML (10MG/ML)</u>	<u>A209967 002</u>	Jan 16, 2020
<u>AP1</u>		<u>100MG/10ML (10MG/ML)</u>	<u>A209967 003</u>	Jan 16, 2020
<u>AP1</u>	SANDOZ	<u>10MG/ML (10MG/ML)</u>	<u>A208905 001</u>	Jan 31, 2019
<u>AP1</u>		<u>50MG/5ML (10MG/ML)</u>	<u>A208905 002</u>	Jan 31, 2019
<u>AP1</u>		<u>100MG/10ML (10MG/ML)</u>	<u>A208905 003</u>	Jan 31, 2019

VAZCULEP

<u>AP1</u>	+	EXELA PHARMA	<u>10MG/ML (10MG/ML)</u>	<u>N204300 001</u>	Jun 27, 2014
<u>AP1</u>	+		<u>50MG/5ML (10MG/ML)</u>	<u>N204300 002</u>	Jun 27, 2014
<u>AP1</u>	+		<u>100MG/10ML (10MG/ML)</u>	<u>N204300 003</u>	Jun 27, 2014

PHENYLEPHRINE HYDROCHLORIDE

<u>AP2</u>		AVET LIFESCIENCES	<u>10MG/ML (10MG/ML)</u>	<u>A209968 001</u>	Feb 28, 2023
<u>AP2</u>		BE PHARMS	<u>10MG/ML (10MG/ML)</u>	<u>A217521 001</u>	Jun 26, 2023
<u>AP2</u>			<u>50MG/5ML (10MG/ML)</u>	<u>A217521 002</u>	Jun 26, 2023
<u>AP2</u>			<u>100MG/10ML (10MG/ML)</u>	<u>A217521 003</u>	Jun 26, 2023
<u>AP2</u>		EUGIA PHARMA	<u>10MG/ML (10MG/ML)</u>	<u>A210696 001</u>	Jan 07, 2021
<u>AP2</u>		FRESENIUS KABI USA	<u>10MG/ML (10MG/ML)</u>	<u>A210665 001</u>	Jan 29, 2019
<u>AP2</u>	+	HIKMA	<u>10MG/ML (10MG/ML)</u>	<u>N203826 001</u>	Dec 20, 2012
<u>AP2</u>		MEITHEAL	<u>10MG/ML (10MG/ML)</u>	<u>A210334 001</u>	Apr 27, 2018

BIORPHEN

+	DR REDDYS LABS SA	0.5MG/5ML (0.1MG/ML)	N212909 001	Oct 21, 2019
---	-------------------	----------------------	-------------	--------------

IMMPHENTIV

+	HIKMA	0.5MG/5ML (0.1MG/ML)	N203826 004	Mar 09, 2023
+		1MG/10ML (0.1MG/ML)	N203826 005	Mar 09, 2023

PHENYLEPHRINE HYDROCHLORIDE

+	HIKMA	50MG/5ML (10MG/ML)	N203826 002	Jun 19, 2019
+		100MG/10ML (10MG/ML)	N203826 003	Jun 19, 2019

SOLUTION/DROPS; OPHTHALMIC

PHENYLEPHRINE HYDROCHLORIDE

<u>AT</u>	+	ALCON	<u>2.5%</u>	<u>N207926 001</u>	Jan 15, 2015
<u>AT</u>	+		<u>10%</u>	<u>N207926 002</u>	Jan 15, 2015
<u>AT</u>		CAPLIN	<u>2.5%</u>	<u>A215183 001</u>	May 23, 2024
<u>AT</u>			<u>10%</u>	<u>A215183 002</u>	May 23, 2024
<u>AT</u>		GLAND PHARMA LTD	<u>2.5%</u>	<u>A218129 001</u>	Dec 18, 2024
<u>AT</u>			<u>10%</u>	<u>A218129 002</u>	Dec 18, 2024
<u>AT</u>	!	MANKIND PHARMA	<u>2.5%</u>	<u>A216859 001</u>	Sep 29, 2022
<u>AT</u>	!		<u>10%</u>	<u>A216496 001</u>	Jan 11, 2023
<u>AT</u>	+	PARAGON BIOTECK	<u>2.5%</u>	<u>N203510 001</u>	Mar 21, 2013
<u>AT</u>	+		<u>10%</u>	<u>N203510 002</u>	Mar 21, 2013

PHENYLEPHRINE HYDROCHLORIDE; PROMETHAZINE HYDROCHLORIDE

SYRUP; ORAL

PHENYLEPHRINE HYDROCHLORIDE AND PROMETHAZINE HYDROCHLORIDE

<u>AA</u>	!	GENUS	<u>5MG/5ML; 6.25MG/5ML</u>	<u>A040654 001</u>	Dec 07, 2006
<u>AA</u>		AMNEAL PHARMS	<u>5MG/5ML; 6.25MG/5ML</u>	<u>A040902 001</u>	Aug 25, 2009

PHENYTOIN

SUSPENSION; ORAL

DILANTIN-125

<u>AB</u>	+	VIATRIS	<u>125MG/5ML</u>	<u>N008762 001</u>	
-----------	---	---------	------------------	--------------------	--

PHENYTOIN

<u>AB</u>		TARO	<u>125MG/5ML</u>	<u>A040521 001</u>	Mar 08, 2004
-----------	--	------	------------------	--------------------	--------------

TABLET, CHEWABLE; ORAL

DILANTIN

<u>AB</u>	+	PHARMACIA	<u>50MG</u>	<u>A084427 001</u>	
-----------	---	-----------	-------------	--------------------	--

PRESCRIPTION DRUG PRODUCT LIST

PHENYTOIN

TABLET, CHEWABLE; ORAL

PHENYTOIN

<u>AB</u>	EPIC PHARMA LLC	<u>50MG</u>	<u>A040884</u>	<u>001</u>	Nov 28, 2014
<u>AB</u>	RISING	<u>50MG</u>	<u>A200691</u>	<u>001</u>	Dec 26, 2012
<u>AB</u>	TARO	<u>50MG</u>	<u>A200565</u>	<u>001</u>	Apr 17, 2014

PHENYTOIN SODIUM

CAPSULE; ORAL

DILANTIN

<u>AB</u>	+! VIATRIS	<u>100MG EXTENDED</u>	<u>A084349</u>	<u>002</u>	
<u>EXTENDED PHENYTOIN SODIUM</u>					
<u>AB</u>	AMNEAL PHARMS NY	<u>100MG EXTENDED</u>	<u>A040765</u>	<u>001</u>	Nov 12, 2008
<u>AB</u>	TARO	<u>100MG EXTENDED</u>	<u>A040684</u>	<u>001</u>	Sep 05, 2006

PHENYTOIN SODIUM

<u>AB</u>	AUROBINDO PHARMA	<u>100MG EXTENDED</u>	<u>A204309</u>	<u>001</u>	Jun 10, 2015
DILANTIN					
	+! VIATRIS	30MG EXTENDED	A084349	001	
PHENYTEK					
	MYLAN	200MG EXTENDED	A040298	002	Dec 06, 2001
	!	300MG EXTENDED	A040298	003	Dec 06, 2001

INJECTABLE; INJECTION

PHENYTOIN SODIUM

<u>AP</u>	ACELLA	<u>50MG/ML</u>	<u>A040573</u>	<u>001</u>	Sep 13, 2006
<u>AP</u>	+! HIKMA	<u>50MG/ML</u>	<u>A084307</u>	<u>001</u>	

PHYTONADIONE

INJECTABLE; INJECTION

PHYTONADIONE

<u>AB</u>	CIPLA	<u>1MG/0.5ML</u>	<u>A214966</u>	<u>001</u>	Jan 03, 2025
<u>AB</u>	+! INTL MEDICATION	<u>1MG/0.5ML</u>	<u>A083722</u>	<u>001</u>	
<u>AB1</u>	DR REDDYS	<u>10MG/ML</u>	<u>A207719</u>	<u>001</u>	May 22, 2019
<u>AB1</u>	GLAND PHARMA LTD	<u>10MG/ML</u>	<u>A217386</u>	<u>001</u>	Dec 10, 2024

VITAMIN K1

<u>AB1</u>	+! HOSPIRA	<u>10MG/ML</u>	<u>A087955</u>	<u>001</u>	Jul 25, 1983
BP	+!	1MG/0.5ML	A087954	001	Jul 25, 1983
PHYTONADIONE					
	! CIPLA	1MG/0.5ML	A214596	001	Apr 22, 2022
	!	10MG/ML	A214596	002	Apr 22, 2022

TABLET; ORAL

PHYTONADIONE

<u>AB</u>	AGNITIO	<u>5MG</u>	<u>A213336</u>	<u>001</u>	Oct 12, 2023
<u>AB</u>	AMNEAL PHARMS CO	<u>5MG</u>	<u>A209373</u>	<u>001</u>	May 11, 2018
<u>AB</u>	HIKMA	<u>5MG</u>	<u>A212112</u>	<u>001</u>	Dec 19, 2024
<u>AB</u>	SCIEGEN PHARMS INC	<u>5MG</u>	<u>A213329</u>	<u>001</u>	Jul 28, 2023
<u>AB</u>	! ZYDUS LIFESCIENCES	<u>5MG</u>	<u>A210189</u>	<u>001</u>	Feb 20, 2019

PIFLUFOLASTAT F-18

SOLUTION; INTRAVENOUS

PYLARIFY

	+! PROGENICS PHARMS INC	50ML (1-80mCi/ML)	N214793	001	May 26, 2021
--	-------------------------	-------------------	---------	-----	--------------

PILOCARPINE HYDROCHLORIDE

SOLUTION; OPHTHALMIC

ISOPTO CARPINE

<u>AT</u>	+! SANDOZ	<u>1%</u>	<u>N200890</u>	<u>001</u>	Jun 22, 2010
<u>AT</u>	+!	<u>2%</u>	<u>N200890</u>	<u>002</u>	Jun 22, 2010
<u>AT</u>	+!	<u>4%</u>	<u>N200890</u>	<u>003</u>	Jun 22, 2010

PILOCARPINE HYDROCHLORIDE

<u>AT</u>	AMNEAL	<u>1%</u>	<u>A214193</u>	<u>001</u>	Sep 21, 2020
<u>AT</u>		<u>2%</u>	<u>A214193</u>	<u>002</u>	Sep 21, 2020
<u>AT</u>		<u>4%</u>	<u>A214193</u>	<u>003</u>	Sep 21, 2020
<u>AT</u>	RISING	<u>1%</u>	<u>A204398</u>	<u>001</u>	Sep 27, 2017
<u>AT</u>		<u>2%</u>	<u>A204398</u>	<u>002</u>	Sep 27, 2017
<u>AT</u>		<u>4%</u>	<u>A204398</u>	<u>003</u>	Sep 27, 2017
<u>AT</u>	SOMERSET THERAPS LLC	<u>1%</u>	<u>A210384</u>	<u>001</u>	Nov 25, 2019
<u>AT</u>		<u>2%</u>	<u>A210384</u>	<u>002</u>	Nov 25, 2019
<u>AT</u>		<u>4%</u>	<u>A210384</u>	<u>003</u>	Nov 25, 2019

QLOSI

	+! ORASIS PHARMS	0.4%	N217836	001	Oct 17, 2023
--	------------------	------	---------	-----	--------------

VUITY

	+! ABBVIE	1.25%	N214028	001	Oct 28, 2021
--	-----------	-------	---------	-----	--------------

PRESCRIPTION DRUG PRODUCT LIST

PILOCARPINE HYDROCHLORIDE

TABLET; ORAL

PILOCARPINE HYDROCHLORIDE

AB	AUROBINDO PHARMA LTD	5MG	A212377 001	Aug 13, 2019
AB		7.5MG	A212377 002	Aug 13, 2019
AB	IMPAX LABS	5MG	A077248 001	Mar 31, 2006
AB		7.5MG	A077248 002	Mar 31, 2006
AB	INNOGENIX	5MG	A076963 001	Dec 22, 2004
AB		7.5MG	A076963 002	Feb 27, 2007
AB	LANNETT CO INC	5MG	A077220 001	Oct 14, 2005
AB		7.5MG	A077220 002	May 06, 2009
AB	PADAGIS US	5MG	A076746 001	Nov 16, 2004
<u>SALAGEN</u>				
AB	+ ADVANZ PHARMA	5MG	N020237 001	Mar 22, 1994
AB	+!	7.5MG	N020237 002	Apr 18, 2003

PIMAVANSERIN TARTRATE

CAPSULE; ORAL

NUPLAZID

+! ACADIA PHARMS INC EQ 34MG BASE N210793 001 Jun 28, 2018

TABLET; ORAL

NUPLAZID

+! ACADIA PHARMS INC EQ 10MG BASE N207318 002 Jun 28, 2018

PIMECROLIMUS

CREAM; TOPICAL

ELIDEL**AB** +! BAUSCH **1%** **N021302 001** Dec 13, 2001**PIMECROLIMUS****AB** ACTAVIS LABS UT INC **1%** **A209345 001** Dec 27, 2018**AB** GLENMARK PHARMS LTD **1%** **A211769 001** Aug 29, 2019PIMOZIDE

TABLET; ORAL

PIMOZIDE

ENDO OPERATIONS 1MG A204521 001 Sep 28, 2015

! 2MG A204521 002 Sep 28, 2015

PINDOLOL

TABLET; ORAL

PINDOLOL**AB** ANI PHARMS **5MG** **A073609 002** Mar 29, 1993**AB** **10MG** **A073609 001** Mar 29, 1993**AB** AUROBINDO PHARMA USA **5MG** **A074019 001** Sep 03, 1992**AB** ! **10MG** **A074019 002** Sep 03, 1992**AB** NOSTRUM LABS INC **5MG** **A205415 001** Jan 13, 2016**AB** **10MG** **A205415 002** Jan 13, 2016**AB** SUN PHARM INDUSTRIES **5MG** **A074063 001** Jan 27, 1994**AB** **10MG** **A074063 002** Jan 27, 1994**AB** UNICHEM **5MG** **A211712 001** Aug 01, 2019**AB** **10MG** **A211712 002** Aug 01, 2019PIOGLITAZONE HYDROCHLORIDE

TABLET; ORAL

ACTOS**AB** + TAKEDA PHARMS USA **EQ 15MG BASE** **N021073 001** Jul 15, 1999**AB** + **EQ 30MG BASE** **N021073 002** Jul 15, 1999**AB** +! **EQ 45MG BASE** **N021073 003** Jul 15, 1999**PIOGLITAZONE HYDROCHLORIDE****AB** ACCORD HLTHCARE **EQ 15MG BASE** **A200044 001** Feb 13, 2013**AB** **EQ 30MG BASE** **A200044 002** Feb 13, 2013**AB** **EQ 45MG BASE** **A200044 003** Feb 13, 2013**AB** ANNORA PHARMA **EQ 15MG BASE** **A204133 001** Apr 07, 2014**AB** **EQ 30MG BASE** **A204133 002** Apr 07, 2014**AB** **EQ 45MG BASE** **A204133 003** Apr 07, 2014**AB** AUROBINDO PHARMA LTD **EQ 15MG BASE** **A200268 001** Feb 13, 2013**AB** **EQ 30MG BASE** **A200268 002** Feb 13, 2013**AB** **EQ 45MG BASE** **A200268 003** Feb 13, 2013**AB** CHARTWELL RX **EQ 15MG BASE** **A076798 001** Oct 26, 2012**AB** **EQ 30MG BASE** **A076798 002** Oct 26, 2012**AB** **EQ 45MG BASE** **A076798 003** Oct 26, 2012**AB** COREPHARMA **EQ 15MG BASE** **A210165 001** Jan 22, 2021

PRESCRIPTION DRUG PRODUCT LIST

PIOGLITAZONE HYDROCHLORIDE

TABLET; ORAL

PIOGLITAZONE HYDROCHLORIDE

<u>AB</u>		<u>EQ 30MG BASE</u>	<u>A210165 002</u>	Jan 22, 2021
<u>AB</u>		<u>EQ 45MG BASE</u>	<u>A210165 003</u>	Jan 22, 2021
<u>AB</u>	MACLEODS PHARMS LTD	<u>EQ 15MG BASE</u>	<u>A202467 001</u>	Feb 06, 2013
<u>AB</u>		<u>EQ 30MG BASE</u>	<u>A202467 002</u>	Feb 06, 2013
<u>AB</u>		<u>EQ 45MG BASE</u>	<u>A202467 003</u>	Feb 06, 2013
<u>AB</u>	PRINSTON INC	<u>EQ 15MG BASE</u>	<u>A207806 001</u>	Apr 17, 2018
<u>AB</u>		<u>EQ 30MG BASE</u>	<u>A207806 002</u>	Apr 17, 2018
<u>AB</u>		<u>EQ 45MG BASE</u>	<u>A207806 003</u>	Apr 17, 2018
<u>AB</u>	PURACAP PHARM LLC	<u>EQ 15MG BASE</u>	<u>A206738 001</u>	Oct 06, 2017
<u>AB</u>		<u>EQ 30MG BASE</u>	<u>A206738 002</u>	Oct 06, 2017
<u>AB</u>		<u>EQ 45MG BASE</u>	<u>A206738 003</u>	Oct 06, 2017
<u>AB</u>	SANDOZ	<u>EQ 15MG BASE</u>	<u>A078670 001</u>	Feb 13, 2013
<u>AB</u>		<u>EQ 30MG BASE</u>	<u>A078670 002</u>	Feb 13, 2013
<u>AB</u>		<u>EQ 45MG BASE</u>	<u>A078670 003</u>	Feb 13, 2013
<u>AB</u>	TEVA PHARMS USA	<u>EQ 15MG BASE</u>	<u>A077210 001</u>	Jan 10, 2014
<u>AB</u>		<u>EQ 30MG BASE</u>	<u>A077210 002</u>	Jan 10, 2014
<u>AB</u>		<u>EQ 45MG BASE</u>	<u>A077210 003</u>	Jan 10, 2014
<u>AB</u>	ZYDUS PHARMS USA INC	<u>EQ 15MG BASE</u>	<u>A202456 001</u>	Feb 13, 2013
<u>AB</u>		<u>EQ 30MG BASE</u>	<u>A202456 002</u>	Feb 13, 2013
<u>AB</u>		<u>EQ 45MG BASE</u>	<u>A202456 003</u>	Feb 13, 2013

PIPERACILLIN SODIUM

INJECTABLE; INJECTION

PIPERACILLIN

!	ISTITUTO BIO ITA SPA	EQ 2GM BASE/VIAL	A065114 001	Nov 14, 2003
!		EQ 3GM BASE/VIAL	A065114 002	Nov 14, 2003
!		EQ 4GM BASE/VIAL	A065114 003	Nov 14, 2003
!		EQ 40GM BASE/VIAL	A065157 001	Jul 12, 2004

PIPERACILLIN SODIUM; TAZOBACTAM SODIUM

INJECTABLE; INJECTION

PIPERACILLIN AND TAZOBACTAM

<u>AP</u>	EUGIA PHARMA SPECLTS	<u>EQ 2GM BASE/VIAL;EQ 250MG BASE/VIAL</u>	<u>A065498 001</u>	May 23, 2011
<u>AP</u>		<u>EQ 3GM BASE/VIAL;EQ 375MG BASE/VIAL</u>	<u>A065498 002</u>	May 23, 2011
<u>AP</u>		<u>EQ 4GM BASE/VIAL;EQ 500MG BASE/VIAL</u>	<u>A065498 003</u>	May 23, 2011
<u>AP</u>	FRESENIUS KABI	<u>EQ 2GM BASE/VIAL;EQ 250MG BASE/VIAL</u>	<u>A203719 001</u>	May 18, 2018
<u>AP</u>		<u>EQ 3GM BASE/VIAL;EQ 375MG BASE/VIAL</u>	<u>A203719 002</u>	May 18, 2018
<u>AP</u>		<u>EQ 4GM BASE/VIAL;EQ 500MG BASE/VIAL</u>	<u>A203719 003</u>	May 18, 2018
<u>AP</u>		<u>EQ 36GM BASE/VIAL;EQ 4.5GM BASE/VIAL</u>	<u>A203720 001</u>	May 11, 2018
<u>AP</u>	ISTITUTO BIO ITA SPA	<u>EQ 2GM BASE/VIAL;EQ 250MG BASE/VIAL</u>	<u>A065523 001</u>	May 31, 2011
<u>AP</u>		<u>EQ 3GM BASE/VIAL;EQ 375MG BASE/VIAL</u>	<u>A065523 002</u>	May 31, 2011
<u>AP</u>		<u>EQ 4GM BASE/VIAL;EQ 500MG BASE/VIAL</u>	<u>A065523 003</u>	May 31, 2011
<u>AP</u>		<u>EQ 36GM BASE/VIAL;EQ 4.5GM BASE/VIAL</u>	<u>A090498 001</u>	May 31, 2011
<u>AP</u>	PROVEPHARM SAS	<u>EQ 2GM BASE/VIAL;EQ 250MG BASE/VIAL</u>	<u>A207847 001</u>	Jan 13, 2017
<u>AP</u>		<u>EQ 3GM BASE/VIAL;EQ 375MG BASE/VIAL</u>	<u>A207847 002</u>	Jan 13, 2017
<u>AP</u>		<u>EQ 36GM BASE/VIAL;EQ 4.5GM BASE/VIAL</u>	<u>A207848 002</u>	Jan 13, 2017
<u>AP</u>		<u>EQ 4GM BASE/VIAL;EQ 500MG BASE/VIAL</u>	<u>A207847 003</u>	Jan 13, 2017
<u>AP</u>		<u>EQ 12GM BASE/VIAL;EQ 1.5GM BASE/VIAL</u>	<u>A207848 001</u>	May 11, 2018
<u>AP</u>	SAGENT PHARMS INC	<u>EQ 2GM BASE/VIAL;EQ 250MG BASE/VIAL</u>	<u>A208674 001</u>	Feb 16, 2021
<u>AP</u>		<u>EQ 3GM BASE/VIAL;EQ 375MG BASE/VIAL</u>	<u>A208674 002</u>	Feb 16, 2021
<u>AP</u>		<u>EQ 4GM BASE/VIAL;EQ 500MG BASE/VIAL</u>	<u>A208674 003</u>	Feb 16, 2021
<u>AP</u>		<u>EQ 36GM BASE/VIAL;EQ 4.5GM BASE/VIAL</u>	<u>A208675 001</u>	Feb 16, 2021
<u>AP</u>	SANDOZ	<u>EQ 2GM BASE/VIAL;EQ 250MG BASE/VIAL</u>	<u>A065362 001</u>	Oct 21, 2010
<u>AP</u>		<u>EQ 2GM BASE/VIAL;EQ 250MG BASE/VIAL</u>	<u>A065363 001</u>	Oct 21, 2010
<u>AP</u>		<u>EQ 3GM BASE/VIAL;EQ 375MG BASE/VIAL</u>	<u>A065362 002</u>	Oct 21, 2010
<u>AP</u>		<u>EQ 3GM BASE/VIAL;EQ 375MG BASE/VIAL</u>	<u>A065363 002</u>	Oct 21, 2010
<u>AP</u>		<u>EQ 4GM BASE/VIAL;EQ 500MG BASE/VIAL</u>	<u>A065362 003</u>	Oct 21, 2010
<u>AP</u>		<u>EQ 4GM BASE/VIAL;EQ 500MG BASE/VIAL</u>	<u>A065363 003</u>	Oct 21, 2010
<u>AP</u>	!	<u>EQ 12GM BASE/VIAL;EQ 1.5GM BASE/VIAL</u>	<u>A203557 001</u>	Oct 29, 2014
<u>AP</u>		<u>EQ 36GM BASE/VIAL;EQ 4.5GM BASE/VIAL</u>	<u>A203557 002</u>	Jul 08, 2021
<u>AP</u>	SHANDONG	<u>EQ 2GM BASE/VIAL;EQ 250MG BASE/VIAL</u>	<u>A204959 001</u>	Aug 10, 2018
<u>AP</u>		<u>EQ 3GM BASE/VIAL;EQ 375MG BASE/VIAL</u>	<u>A204959 002</u>	Aug 10, 2018
<u>AP</u>		<u>EQ 4GM BASE/VIAL;EQ 500MG BASE/VIAL</u>	<u>A204959 003</u>	Aug 10, 2018
<u>AP</u>		<u>EQ 36GM BASE/VIAL;EQ 4.5GM BASE/VIAL</u>	<u>A217243 001</u>	Apr 14, 2023
<u>AP</u>	STERISCIENCE SPECLTS	<u>EQ 2GM BASE/VIAL;EQ 250MG BASE/VIAL</u>	<u>A065458 001</u>	Aug 15, 2014
<u>AP</u>		<u>EQ 3GM BASE/VIAL;EQ 375MG BASE/VIAL</u>	<u>A065458 002</u>	Aug 15, 2014

PRESCRIPTION DRUG PRODUCT LIST

PIPERACILLIN SODIUM; TAZOBACTAM SODIUM

INJECTABLE; INJECTION

PIPERACILLIN AND TAZOBACTAM

<u>AP</u>		<u>EQ 4GM BASE/VIAL;EQ 500MG BASE/VIAL</u>	<u>A065458 003</u>	Aug 15, 2014
<u>AP</u>	!	WOCKHARDT BIO AG <u>EQ 2GM BASE/VIAL;EQ 250MG BASE/VIAL</u>	<u>A206996 001</u>	Mar 22, 2017
<u>AP</u>	!	<u>EQ 3GM BASE/VIAL;EQ 375MG BASE/VIAL</u>	<u>A206996 002</u>	Mar 22, 2017
<u>AP</u>	!	<u>EQ 4GM BASE/VIAL;EQ 500MG BASE/VIAL</u>	<u>A206996 003</u>	Mar 22, 2017
<u>AP</u>	!	<u>EQ 36GM BASE/VIAL;EQ 4.5GM BASE/VIAL</u>	<u>A207146 001</u>	Mar 17, 2017
ZOSYN IN PLASTIC CONTAINER				
	+	!	BAXTER HLTHCARE CORP	EQ 40MG BASE/ML;EQ 5MG BASE/ML N050750 001 Feb 24, 1998
	+	!		EQ 60MG BASE/ML;EQ 7.5MG BASE/ML N050750 002 Feb 24, 1998
	+	!		EQ 4GM BASE/100ML;EQ 500MG BASE/100ML N050750 003 Feb 24, 1998

PIRFENIDONE

CAPSULE; ORAL

ESBRIET

<u>AB</u>	+	!	GENENTECH INC	<u>267MG</u>	<u>N022535 001</u>	Oct 15, 2014
-----------	---	---	---------------	--------------	--------------------	--------------

PIRFENIDONE

<u>AB</u>			ACCORD HLTHCARE	<u>267MG</u>	<u>A212731 001</u>	Jan 20, 2022
<u>AB</u>			AMNEAL	<u>267MG</u>	<u>A212569 001</u>	Jan 03, 2022
<u>AB</u>			LAURUS	<u>267MG</u>	<u>A212724 001</u>	Jul 19, 2022
<u>AB</u>			MACLEODS PHARMS LTD	<u>267MG</u>	<u>A212748 001</u>	Dec 18, 2024
<u>AB</u>			SANDOZ	<u>267MG</u>	<u>A212600 001</u>	Jun 13, 2022
<u>AB</u>			SCIEGEN PHARMS INC	<u>267MG</u>	<u>A212077 001</u>	Aug 01, 2022

TABLET; ORAL

ESBRIET

<u>AB</u>	+		GENENTECH INC	<u>267MG</u>	<u>N208780 001</u>	Jan 11, 2017
<u>AB</u>	+	!		<u>801MG</u>	<u>N208780 003</u>	Jan 11, 2017

PIRFENIDONE

<u>AB</u>			ACCORD HLTHCARE	<u>267MG</u>	<u>A212730 001</u>	Jan 25, 2022
<u>AB</u>				<u>801MG</u>	<u>A212730 002</u>	Jan 25, 2022
<u>AB</u>			AIZANT	<u>267MG</u>	<u>A212747 001</u>	Jul 21, 2022
<u>AB</u>				<u>534MG</u>	<u>A212747 002</u>	Jul 21, 2022
<u>AB</u>				<u>801MG</u>	<u>A212747 003</u>	Jul 21, 2022
<u>AB</u>			ALEMBIC	<u>267MG</u>	<u>A212708 001</u>	May 20, 2022
<u>AB</u>				<u>801MG</u>	<u>A212708 002</u>	May 20, 2022
<u>AB</u>			AMNEAL	<u>267MG</u>	<u>A212570 001</u>	Mar 25, 2022
<u>AB</u>				<u>801MG</u>	<u>A212570 002</u>	Mar 25, 2022
<u>AB</u>			APOTEX	<u>267MG</u>	<u>A212709 001</u>	May 31, 2023
<u>AB</u>				<u>801MG</u>	<u>A212709 002</u>	May 31, 2023
<u>AB</u>			HETERO LABS LTD V	<u>267MG</u>	<u>A212674 001</u>	Sep 21, 2022
<u>AB</u>				<u>801MG</u>	<u>A212674 002</u>	Sep 21, 2022
<u>AB</u>			LAURUS	<u>267MG</u>	<u>A212722 001</u>	Jul 19, 2022
<u>AB</u>				<u>534MG</u>	<u>A212722 002</u>	Jul 19, 2022
<u>AB</u>				<u>801MG</u>	<u>A212722 003</u>	Jul 19, 2022
<u>AB</u>			MICRO LABS	<u>267MG</u>	<u>A212680 001</u>	May 18, 2022
<u>AB</u>				<u>801MG</u>	<u>A212680 002</u>	May 18, 2022
<u>AB</u>			MSN	<u>267MG</u>	<u>A212772 001</u>	May 24, 2022
<u>AB</u>				<u>801MG</u>	<u>A212772 002</u>	May 24, 2022
<u>AB</u>			SANDOZ	<u>267MG</u>	<u>A212560 001</u>	Apr 28, 2022
<u>AB</u>				<u>801MG</u>	<u>A212560 002</u>	Apr 28, 2022
<u>AB</u>			SCIEGEN PHARMS INC	<u>267MG</u>	<u>A212078 001</u>	Aug 01, 2022
<u>AB</u>				<u>801MG</u>	<u>A212078 002</u>	Aug 01, 2022
<u>AB</u>			TEVA PHARMS USA	<u>267MG</u>	<u>A212759 001</u>	Jan 25, 2022
<u>AB</u>				<u>801MG</u>	<u>A212759 002</u>	Jan 25, 2022

PIROXICAM

CAPSULE; ORAL

FELDENE

<u>AB</u>	+		PFIZER	<u>10MG</u>	<u>N018147 002</u>	Apr 06, 1982
<u>AB</u>	+	!		<u>20MG</u>	<u>N018147 003</u>	Apr 06, 1982

PIROXICAM

<u>AB</u>			MICRO LABS	<u>10MG</u>	<u>A206152 001</u>	Dec 29, 2017
<u>AB</u>				<u>20MG</u>	<u>A206152 002</u>	Dec 29, 2017
<u>AB</u>			NOSTRUM LABS INC	<u>10MG</u>	<u>A074118 002</u>	Jun 15, 1993
<u>AB</u>				<u>20MG</u>	<u>A074118 001</u>	Jun 15, 1993
<u>AB</u>			STRIDES PHARMA	<u>10MG</u>	<u>A206136 001</u>	Jun 20, 2017
<u>AB</u>				<u>10MG</u>	<u>A210347 001</u>	Jan 26, 2018
<u>AB</u>				<u>20MG</u>	<u>A206136 002</u>	Jun 20, 2017
<u>AB</u>				<u>20MG</u>	<u>A210347 002</u>	Jan 26, 2018
<u>AB</u>			TEVA	<u>10MG</u>	<u>A074131 001</u>	Dec 11, 1992
<u>AB</u>				<u>20MG</u>	<u>A074131 002</u>	Dec 11, 1992
<u>AB</u>			UNICHEM	<u>10MG</u>	<u>A208340 001</u>	Apr 13, 2017

PRESCRIPTION DRUG PRODUCT LIST

PIROXICAM

CAPSULE;ORAL

PIROXICAM

AB		20MG	A208340 002	Apr 13, 2017
AB	ZYDUS LIFESCIENCES	10MG	A205585 001	Jul 17, 2018
AB		20MG	A205585 002	Jul 17, 2018

PIRTOBRUTINIB

TABLET;ORAL

JAYPIRCA

+	LOXO ONCOL	50MG	N216059 001	Jan 27, 2023
+	!	100MG	N216059 002	Jan 27, 2023

PITAVASTATIN CALCIUM

TABLET;ORAL

LIVALO

AB	+	KOWA CO	EQ 1MG BASE	N022363 001	Aug 03, 2009
AB	+		EQ 2MG BASE	N022363 002	Aug 03, 2009
AB	+	!	EQ 4MG BASE	N022363 003	Aug 03, 2009

PITAVASTATIN CALCIUM

AB		AMNEAL PHARMS NY	EQ 1MG BASE	A205961 001	Aug 20, 2024
AB			EQ 2MG BASE	A205961 002	Aug 20, 2024
AB			EQ 4MG BASE	A205961 003	Aug 20, 2024
AB		AUROBINDO PHARMA	EQ 1MG BASE	A206015 001	Dec 20, 2016
AB			EQ 2MG BASE	A206015 002	Dec 20, 2016
AB			EQ 4MG BASE	A206015 003	Dec 20, 2016
AB		HETERO LABS LTD V	EQ 1MG BASE	A205977 001	Apr 30, 2024
AB			EQ 2MG BASE	A205977 002	Apr 30, 2024
AB			EQ 4MG BASE	A205977 003	Apr 30, 2024
AB		LUPIN LTD	EQ 1MG BASE	A206029 001	Nov 20, 2023
AB			EQ 2MG BASE	A206029 002	Nov 20, 2023
AB			EQ 4MG BASE	A206029 003	Nov 20, 2023
AB		MYLAN	EQ 1MG BASE	A206070 001	Apr 04, 2019
AB			EQ 2MG BASE	A206070 002	Apr 04, 2019
AB			EQ 4MG BASE	A206070 003	Apr 04, 2019
AB		ORIENT PHARMA CO LTD	EQ 1MG BASE	A205932 001	Feb 03, 2017
AB			EQ 2MG BASE	A205932 002	Feb 03, 2017
AB			EQ 4MG BASE	A205932 003	Feb 03, 2017
AB		SAWAI USA	EQ 1MG BASE	A205955 001	Feb 03, 2017
AB			EQ 2MG BASE	A205955 002	Feb 03, 2017
AB			EQ 4MG BASE	A205955 003	Feb 03, 2017
AB		ZYDUS PHARMS	EQ 1MG BASE	A206047 001	Feb 22, 2023
AB			EQ 2MG BASE	A206047 002	Feb 22, 2023
AB			EQ 4MG BASE	A206047 003	Feb 22, 2023

PITAVASTATIN MAGNESIUM

TABLET;ORAL

ZYPITAMAG

+	MEDICURE	EQ 2MG BASE	N208379 002	Jul 14, 2017
+	!	EQ 4MG BASE	N208379 003	Jul 14, 2017

PITOLISANT HYDROCHLORIDE

TABLET;ORAL

WAKIX

+	HARMONY	EQ 4.45MG BASE	N211150 001	Aug 14, 2019
+	!	EQ 17.8MG BASE	N211150 002	Aug 14, 2019

PIVMECILLINAM HYDROCHLORIDE

TABLET;ORAL

PIVYA

+	UTILITY THERAP	EQ 185MG BASE	N216483 001	Apr 24, 2024
---	----------------	---------------	-------------	--------------

PLAZOMICIN SULFATE

SOLUTION;INTRAVENOUS

ZEMDRI

+	CIPLA USA	EQ 500MG BASE/10ML (EQ 50MG BASE/ML)	N210303 001	Jun 25, 2018
---	-----------	--------------------------------------	-------------	--------------

PLECANATIDE

TABLET;ORAL

TRULANCE

+	SALIX	3MG	N208745 001	Jan 19, 2017
---	-------	-----	-------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

PLERIXAFOR

SOLUTION;SUBCUTANEOUS

MOZOBI

AP	+ !	GENZYME	<u>24MG/1.2ML (20MG/ML)</u>	<u>N022311</u>	<u>001</u>	Dec 15, 2008
-----------	------------	---------	-----------------------------	----------------	------------	--------------

PLERIXAFOR

AP		AMNEAL	<u>24MG/1.2ML (20MG/ML)</u>	<u>A215334</u>	<u>001</u>	Jul 24, 2023
AP		DR REDDYS	<u>24MG/1.2ML (20MG/ML)</u>	<u>A205182</u>	<u>001</u>	Jul 24, 2023
AP		EUGIA PHARMA	<u>24MG/1.2ML (20MG/ML)</u>	<u>A213672</u>	<u>001</u>	Jul 24, 2023
AP		GLAND PHARMA LTD	<u>24MG/1.2ML (20MG/ML)</u>	<u>A206644</u>	<u>001</u>	May 03, 2024
AP		MEITHEAL	<u>24MG/1.2ML (20MG/ML)</u>	<u>A215698</u>	<u>001</u>	Jul 24, 2023
AP		MSN	<u>24MG/1.2ML (20MG/ML)</u>	<u>A211901</u>	<u>001</u>	Jul 24, 2023
AP		TEVA PHARMS USA INC	<u>24MG/1.2ML (20MG/ML)</u>	<u>A205197</u>	<u>001</u>	Jul 24, 2023

PODOFILOX

GEL;TOPICAL

CONDYLOX

AB	+ !	ALLERGAN	<u>0.5%</u>	<u>N020529</u>	<u>001</u>	Mar 13, 1997
-----------	------------	----------	-------------	----------------	------------	--------------

PODOFILOX

AB		PADAGIS US	<u>0.5%</u>	<u>A211871</u>	<u>001</u>	Nov 22, 2023
-----------	--	------------	-------------	----------------	------------	--------------

SOLUTION;TOPICAL

CONDYLOX

AT	+ !	TEVA BRANDED PHARM	<u>0.5%</u>	<u>N019795</u>	<u>001</u>	Dec 13, 1990
-----------	------------	--------------------	-------------	----------------	------------	--------------

PODOFILOX

AT		PADAGIS US	<u>0.5%</u>	<u>A075600</u>	<u>001</u>	Jan 29, 2002
-----------	--	------------	-------------	----------------	------------	--------------

POLIDOCANOL

SOLUTION;INTRAVENOUS

ASCLERA

	+	CHEMISCH FBRK	10MG/2ML (5MG/ML)	N021201	001	Mar 30, 2010
	+ !	KRSSLR	20MG/2ML (10MG/ML)	N021201	002	Mar 30, 2010

VARITHENA

	+ !	PROVENSIS	180MG/18ML (10MG/ML)	N205098	001	Nov 25, 2013
--	------------	-----------	----------------------	---------	-----	--------------

POLYETHYLENE GLYCOL 3350; POTASSIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE

FOR SOLUTION;ORAL

NULYTELY

AA	+ !	BRAINTREE	<u>420GM/BOT;1.48GM/BOT;5.72GM/BOT;11.2GM/BOT</u>	<u>N019797</u>	<u>001</u>	Apr 22, 1991
-----------	------------	-----------	---	----------------	------------	--------------

NULYTELY-FLAVORED

AA	+ !	BRAINTREE	<u>420GM/BOT;1.48GM/BOT;5.72GM/BOT;11.2GM/BOT</u>	<u>N019797</u>	<u>002</u>	Nov 18, 1994
-----------	------------	-----------	---	----------------	------------	--------------

PEG-3350, POTASSIUM CHLORIDE, SODIUM BICARBONATE, SODIUM CHLORIDE

AA		NOVEL LABS INC	<u>420GM/BOT;1.48GM/BOT;5.72GM/BOT;11.2GM/BOT</u>	<u>A090019</u>	<u>001</u>	May 27, 2009
-----------	--	----------------	---	----------------	------------	--------------

AA		STRIDES PHARMA	<u>420GM/BOT;1.48GM/BOT;5.72GM/BOT;11.2GM/BOT</u>	<u>A204559</u>	<u>001</u>	Apr 13, 2015
-----------	--	----------------	---	----------------	------------	--------------

POLYETHYLENE GLYCOL 3350; POTASSIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE; SODIUM SULFATE ANHYDROUS

FOR SOLUTION;ORAL

GOLYTELY

AA	+ !	BRAINTREE	<u>236GM/BOT;2.97GM/BOT;6.74GM/BOT;5.86GM/BOT;22.74GM/BOT</u>	<u>N019011</u>	<u>001</u>	Jul 13, 1984
-----------	------------	-----------	---	----------------	------------	--------------

PEG 3350 AND ELECTROLYTES

AA		NOVEL LABS INC	<u>236GM/BOT;2.97GM/BOT;6.74GM/BOT;5.86GM/BOT;22.74GM/BOT</u>	<u>A090231</u>	<u>001</u>	Jun 01, 2009
-----------	--	----------------	---	----------------	------------	--------------

POLYETHYLENE GLYCOL 3350 AND ELECTROLYTES

AA		STRIDES PHARMA	<u>236GM/BOT;2.97GM/BOT;6.74GM/BOT;5.86GM/BOT;22.74GM/BOT</u>	<u>A204558</u>	<u>001</u>	Dec 21, 2018
-----------	--	----------------	---	----------------	------------	--------------

PEG 3350 AND ELECTROLYTES

		NOVEL LABS INC	240GM/BOT;2.98GM/BOT;6.72GM/BOT;5.84GM/BOT;22.72GM/BOT	A090186	001	Jun 01, 2009
--	--	----------------	--	---------	-----	--------------

POLYMYXIN B SULFATE

INJECTABLE;INJECTION

POLYMYXIN B SULFATE

AP		ADRASTEIA PHARMA	<u>EQ 500,000 UNITS BASE/VIAL</u>	<u>A063000</u>	<u>001</u>	Sep 30, 1994
AP		EUGIA PHARMA	<u>EQ 500,000 UNITS BASE/VIAL</u>	<u>A206589</u>	<u>001</u>	Apr 04, 2016
AP		FRESENIUS KABI USA	<u>EQ 500,000 UNITS BASE/VIAL</u>	<u>A065372</u>	<u>001</u>	Jan 10, 2008
AP		GLAND	<u>EQ 500,000 UNITS BASE/VIAL</u>	<u>A207322</u>	<u>001</u>	Apr 14, 2016
AP	!	XELLIA PHARMS APS	<u>EQ 500,000 UNITS BASE/VIAL</u>	<u>A202766</u>	<u>001</u>	Jan 15, 2014

PRESCRIPTION DRUG PRODUCT LIST

POLYMYXIN B SULFATE; TRIMETHOPRIM SULFATE

SOLUTION/DROPS;OPHTHALMIC

TRIMETHOPRIM SULFATE AND POLYMYXIN B SULFATE

<u>AT</u>	BAUSCH AND LOMB	<u>10,000 UNITS/ML;EQ 1MG BASE/ML</u>	<u>A064120 001</u>	Feb 14, 1997
<u>AT</u>	EPIC PHARMA LLC	<u>10,000 UNITS/ML;EQ 1MG BASE/ML</u>	<u>A065006 001</u>	Dec 17, 1998
<u>AT</u>	! SANDOZ	<u>10,000 UNITS/ML;EQ 1MG BASE/ML</u>	<u>A064211 001</u>	Apr 13, 1998
<u>AT</u>	SOMERSET THERAPS LLC	<u>10,000 UNITS/ML;EQ 1MG BASE/ML</u>	<u>A211572 001</u>	Aug 05, 2024

POMALIDOMIDE

CAPSULE;ORAL

POMALIDOMIDE

<u>AB</u>	APOTEX	<u>1MG</u>	<u>A210164 001</u>	Jun 11, 2024
<u>AB</u>		<u>2MG</u>	<u>A210164 002</u>	Jun 11, 2024
<u>AB</u>		<u>3MG</u>	<u>A210164 003</u>	Jun 11, 2024
<u>AB</u>		<u>4MG</u>	<u>A210164 004</u>	Jun 11, 2024
<u>AB</u>	HETERO LABS LTD V	<u>1MG</u>	<u>A210236 001</u>	Sep 26, 2024
<u>AB</u>		<u>2MG</u>	<u>A210236 002</u>	Sep 26, 2024
<u>AB</u>		<u>3MG</u>	<u>A210236 003</u>	Sep 26, 2024
<u>AB</u>		<u>4MG</u>	<u>A210236 004</u>	Sep 26, 2024

POMALYST

<u>AB</u>	+ BRISTOL	<u>1MG</u>	<u>N204026 001</u>	Feb 08, 2013
<u>AB</u>	+	<u>2MG</u>	<u>N204026 002</u>	Feb 08, 2013
<u>AB</u>	+	<u>3MG</u>	<u>N204026 003</u>	Feb 08, 2013
<u>AB</u>	+!	<u>4MG</u>	<u>N204026 004</u>	Feb 08, 2013

PONATINIB HYDROCHLORIDE

TABLET;ORAL

ICLUSIG

+	TAKEDA PHARMS USA	EQ 10MG BASE	N203469 004	Dec 18, 2020
+		EQ 15MG BASE	N203469 001	Dec 14, 2012
+		EQ 30MG BASE	N203469 003	Apr 23, 2015
+	!	EQ 45MG BASE	N203469 002	Dec 14, 2012

PONESIMOD

TABLET;ORAL

PONVORY

+	!	VANDA PHARMS INC	2MG	N213498 001	Mar 18, 2021
+			3MG	N213498 002	Mar 18, 2021
+			4MG	N213498 003	Mar 18, 2021
+			5MG	N213498 004	Mar 18, 2021
+			6MG	N213498 005	Mar 18, 2021
+			7MG	N213498 006	Mar 18, 2021
+			8MG	N213498 007	Mar 18, 2021
+			9MG	N213498 008	Mar 18, 2021
+			10MG	N213498 009	Mar 18, 2021
+			20MG	N213498 010	Mar 18, 2021

PORFIMER SODIUM

INJECTABLE;INJECTION

PHOTOFRIN

	PINNACLE BIOLGS	75MG/VIAL	N020451 001	Dec 27, 1995
--	-----------------	-----------	-------------	--------------

POSACONAZOLE

FOR SUSPENSION, DELAYED RELEASE;ORAL

NOXAFIL POWDERMIX KIT

+	!	MSD MERCK CO	300MG	N214770 001	May 31, 2021
---	---	--------------	-------	-------------	--------------

SOLUTION;INTRAVENOUS

NOXAFIL

<u>AP</u>	+	!	MERCK SHARP DOHME	<u>300MG/16.7ML (18MG/ML)</u>	<u>N205596 001</u>	Mar 13, 2014
-----------	---	---	-------------------	-------------------------------	--------------------	--------------

POSACONAZOLE

<u>AP</u>			ASPIRO	<u>300MG/16.7ML (18MG/ML)</u>	<u>A219057 001</u>	Dec 23, 2024
<u>AP</u>			ENDO OPERATIONS	<u>300MG/16.7ML (18MG/ML)</u>	<u>A208768 001</u>	May 25, 2022
<u>AP</u>			EUGIA PHARMA	<u>300MG/16.7ML (18MG/ML)</u>	<u>A214842 001</u>	Dec 26, 2023
<u>AP</u>			FRESENIUS KABI USA	<u>300MG/16.7ML (18MG/ML)</u>	<u>A209983 001</u>	Dec 26, 2023
<u>AP</u>			GLAND PHARMA LTD	<u>300MG/16.7ML (18MG/ML)</u>	<u>A217553 001</u>	Dec 26, 2023
<u>AP</u>			MYLAN LABS LTD	<u>300MG/16.7ML (18MG/ML)</u>	<u>A211500 001</u>	Dec 26, 2023

SUSPENSION;ORAL

NOXAFIL

<u>AB</u>	+	!	SCHERING	<u>40MG/ML</u>	<u>N022003 001</u>	Sep 15, 2006
-----------	---	---	----------	----------------	--------------------	--------------

POSACONAZOLE

<u>AB</u>			HIKMA	<u>40MG/ML</u>	<u>A208773 001</u>	May 15, 2020
-----------	--	--	-------	----------------	--------------------	--------------

TABLET, DELAYED RELEASE;ORAL

POSACONAZOLE

<u>AB</u>	!		AET PHARMA	<u>100MG</u>	<u>A213454 001</u>	Feb 01, 2021
-----------	---	--	------------	--------------	--------------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

POSACONAZOLE

TABLET, DELAYED RELEASE;ORAL

POSACONAZOLE

<u>AB</u>	AMNEAL	<u>100MG</u>	<u>A216626</u>	<u>001</u>	Dec 22, 2022
<u>AB</u>	AUROBINDO PHARMA	<u>100MG</u>	<u>A217119</u>	<u>001</u>	Jul 18, 2024
<u>AB</u>	BIOCON PHARMA	<u>100MG</u>	<u>A214476</u>	<u>001</u>	Feb 04, 2022
<u>AB</u>	DR REDDYS	<u>100MG</u>	<u>A212500</u>	<u>001</u>	Apr 07, 2022
<u>AB</u>	HETERO LABS LTD III	<u>100MG</u>	<u>A214321</u>	<u>001</u>	Dec 08, 2022
<u>AB</u>	I 3 PHARMS	<u>100MG</u>	<u>A216488</u>	<u>001</u>	Aug 28, 2023
<u>AB</u>	SINOTHERAPEUTICS INC	<u>100MG</u>	<u>A212411</u>	<u>001</u>	Aug 21, 2019
<u>AB</u>	SPECGX LLC	<u>100MG</u>	<u>A212226</u>	<u>001</u>	May 10, 2022
<u>AB</u>	WESTMINSTER PHARMS	<u>100MG</u>	<u>A216326</u>	<u>001</u>	Jun 20, 2023

POTASSIUM ACETATE

INJECTABLE;INJECTION

POTASSIUM ACETATE

<u>AP</u>	EXELA PHARMA	<u>2MEQ/ML</u>	<u>A206203</u>	<u>001</u>	Dec 29, 2015
<u>AP</u>		<u>2MEQ/ML</u>	<u>A212692</u>	<u>001</u>	Oct 20, 2021
<u>AP</u>	FRESENIUS KABI USA	<u>2MEQ/ML</u>	<u>A217515</u>	<u>001</u>	Nov 27, 2024
<u>AP</u>	+! HOSPIRA	<u>2MEQ/ML</u>	<u>N018896</u>	<u>001</u>	Jul 20, 1984

POTASSIUM CHLORIDE

CAPSULE, EXTENDED RELEASE;ORAL

KLOR-CON

<u>AB</u>	UPSHER SMITH LABS	<u>8MEQ</u>	<u>A203106</u>	<u>001</u>	Jul 10, 2015
<u>AB</u>		<u>10MEQ</u>	<u>A203106</u>	<u>002</u>	Jul 10, 2015

POTASSIUM CHLORIDE

<u>AB</u>	ACTAVIS LABS FL INC	<u>8MEQ</u>	<u>A077419</u>	<u>001</u>	Jun 02, 2008
<u>AB</u>	!	<u>10MEQ</u>	<u>A077419</u>	<u>002</u>	Jun 02, 2008
<u>AB</u>	ADARE PHARMS INC	<u>8MEQ</u>	<u>A208864</u>	<u>001</u>	Mar 17, 2017
<u>AB</u>		<u>10MEQ</u>	<u>A208864</u>	<u>002</u>	Mar 17, 2017
<u>AB</u>	AMNEAL PHARMS	<u>10MEQ</u>	<u>A202128</u>	<u>001</u>	Feb 22, 2013
<u>AB</u>	GRANULES	<u>8MEQ</u>	<u>A214686</u>	<u>001</u>	Feb 16, 2021
<u>AB</u>		<u>10MEQ</u>	<u>A214686</u>	<u>002</u>	Feb 16, 2021
<u>AB</u>	LUPIN LTD	<u>8MEQ</u>	<u>A203002</u>	<u>001</u>	Dec 18, 2015
<u>AB</u>		<u>10MEQ</u>	<u>A203002</u>	<u>002</u>	Dec 18, 2015
<u>AB</u>	NOVEL LABS INC	<u>8MEQ</u>	<u>A204828</u>	<u>001</u>	Aug 16, 2016
<u>AB</u>		<u>10MEQ</u>	<u>A204828</u>	<u>002</u>	Aug 16, 2016
<u>AB</u>	PADAGIS US	<u>8MEQ</u>	<u>A200185</u>	<u>001</u>	May 18, 2011
<u>AB</u>		<u>10MEQ</u>	<u>A200185</u>	<u>002</u>	May 18, 2011
<u>AB</u>	PRINSTON INC	<u>8MEQ</u>	<u>A209026</u>	<u>001</u>	Jun 11, 2019
<u>AB</u>		<u>10MEQ</u>	<u>A209026</u>	<u>002</u>	Jun 11, 2019
<u>AB</u>	STRIDES PHARMA	<u>8MEQ</u>	<u>A205549</u>	<u>001</u>	Dec 08, 2015
<u>AB</u>		<u>10MEQ</u>	<u>A205549</u>	<u>002</u>	Dec 08, 2015
<u>AB</u>	ZYDUS PHARMS	<u>8MEQ</u>	<u>A208445</u>	<u>001</u>	Mar 11, 2019
<u>AB</u>		<u>10MEQ</u>	<u>A208445</u>	<u>002</u>	Mar 11, 2019

FOR SOLUTION;ORAL

KLOR-CON

<u>AA</u>	UPSHER SMITH LABS	<u>20MEQ</u>	<u>A209662</u>	<u>001</u>	Oct 23, 2017
-----------	-------------------	--------------	----------------	------------	--------------

POTASSIUM CHLORIDE

<u>AA</u>	AMNEAL	<u>20MEQ</u>	<u>A210902</u>	<u>001</u>	May 23, 2019
<u>AA</u>	BELCHER	<u>20MEQ</u>	<u>A212183</u>	<u>001</u>	May 06, 2019
<u>AA</u>	EPIC PHARMA LLC	<u>20MEQ</u>	<u>A210200</u>	<u>001</u>	Nov 23, 2018
<u>AA</u>	+ GENUS	<u>20MEQ</u>	<u>N208019</u>	<u>001</u>	Aug 19, 2015
<u>AA</u>	GRANULES	<u>20MEQ</u>	<u>A213467</u>	<u>001</u>	Jan 27, 2022
<u>AA</u>	NOVEL LABS INC	<u>20MEQ</u>	<u>A210241</u>	<u>001</u>	Nov 21, 2018
<u>AA</u>	NOVITIUM PHARMA	<u>20MEQ</u>	<u>A212816</u>	<u>001</u>	Jul 12, 2023
<u>AA</u>	RUBICON	<u>20MEQ</u>	<u>A214108</u>	<u>001</u>	Mar 24, 2022
<u>AA</u>	STRIDES PHARMA	<u>20MEQ</u>	<u>A211667</u>	<u>001</u>	Mar 11, 2021

POKONZA

+ GENUS 10MEQ N208019 002 Aug 25, 2023

POTASSIUM CHLORIDE

+! GENUS 40MEQ N208019 003 Aug 25, 2023

INJECTABLE;INJECTION

POTASSIUM CHLORIDE

<u>AP</u>	B BRAUN	<u>2MEQ/ML</u>	<u>A085870</u>	<u>001</u>	
<u>AP</u>	+! HOSPIRA	<u>2MEQ/ML</u>	<u>A080205</u>	<u>001</u>	
<u>AP</u>	NEXUS PHARMS	<u>2MEQ/ML</u>	<u>A217704</u>	<u>001</u>	Aug 14, 2023

POTASSIUM CHLORIDE 10MEQ

<u>AP</u>	FRESENIUS KABI USA	<u>14.9MG/ML</u>	<u>A211087</u>	<u>001</u>	Sep 09, 2020
<u>AP</u>		<u>746MG/100ML</u>	<u>A211087</u>	<u>002</u>	Sep 09, 2020
<u>AP</u>	NEXUS	<u>14.9MG/ML</u>	<u>A214727</u>	<u>001</u>	Mar 18, 2021
<u>AP</u>		<u>745MG/100ML</u>	<u>A214727</u>	<u>002</u>	Mar 18, 2021

PRESCRIPTION DRUG PRODUCT LIST

POTASSIUM CHLORIDE

INJECTABLE; INJECTION

POTASSIUM CHLORIDE 10MEQ IN PLASTIC CONTAINER

<u>AP</u>	+	BAXTER HLTHCARE	<u>14.9MG/ML</u>	<u>N019904</u>	<u>001</u>	Dec 26, 1989
<u>AP</u>	+		<u>746MG/100ML</u>	<u>N019904</u>	<u>005</u>	Dec 17, 1990
<u>AP</u>	+	ICU MEDICAL INC	<u>14.9MG/ML</u>	<u>N020161</u>	<u>005</u>	Nov 30, 1992
<u>AP</u>	+		<u>745MG/100ML</u>	<u>N020161</u>	<u>001</u>	Nov 30, 1992

POTASSIUM CHLORIDE 20MEQ

<u>AP</u>		FRESENIUS KABI USA	<u>29.8MG/ML</u>	<u>A211087</u>	<u>003</u>	Sep 09, 2020
<u>AP</u>			<u>1.49GM/100ML</u>	<u>A211087</u>	<u>005</u>	May 07, 2021
<u>AP</u>		NEXUS	<u>29.8MG/ML</u>	<u>A214727</u>	<u>003</u>	Mar 18, 2021
<u>AP</u>			<u>1.49GM/100ML</u>	<u>A214727</u>	<u>004</u>	Mar 18, 2021

POTASSIUM CHLORIDE 20MEQ IN PLASTIC CONTAINER

<u>AP</u>	+	BAXTER HLTHCARE	<u>29.8MG/ML</u>	<u>N019904</u>	<u>002</u>	Dec 26, 1989
<u>AP</u>	+		<u>1.49GM/100ML</u>	<u>N019904</u>	<u>006</u>	Dec 17, 1990
<u>AP</u>	+	ICU MEDICAL INC	<u>29.8MG/ML</u>	<u>N020161</u>	<u>006</u>	Aug 11, 1998
<u>AP</u>	+		<u>1.49GM/100ML</u>	<u>N020161</u>	<u>002</u>	Nov 30, 1992

POTASSIUM CHLORIDE 40MEQ

<u>AP</u>		FRESENIUS KABI USA	<u>2.98GM/100ML</u>	<u>A211087</u>	<u>004</u>	Sep 09, 2020
<u>AP</u>		NEXUS	<u>2.98GM/100ML</u>	<u>A214727</u>	<u>005</u>	Mar 18, 2021

POTASSIUM CHLORIDE 40MEQ IN PLASTIC CONTAINER

<u>AP</u>	+	BAXTER HLTHCARE	<u>2.98GM/100ML</u>	<u>N019904</u>	<u>004</u>	Dec 26, 1989
<u>AP</u>	+	ICU MEDICAL INC	<u>2.98GM/100ML</u>	<u>N020161</u>	<u>004</u>	Aug 11, 1998

POTASSIUM CHLORIDE IN PLASTIC CONTAINER

<u>AP</u>		FRESENIUS KABI USA	<u>2MEQ/ML</u>	<u>A088901</u>	<u>001</u>	Jan 25, 1985
<u>AP</u>			<u>2MEQ/ML</u>	<u>A088908</u>	<u>001</u>	Jan 25, 1985

POTASSIUM CHLORIDE 30MEQ IN PLASTIC CONTAINER

	+	BAXTER HLTHCARE	2.24GM/100ML	N019904	003	Dec 26, 1989
--	---	-----------------	--------------	---------	-----	--------------

SOLUTION; ORAL

POTASSIUM CHLORIDE

<u>AA</u>		AMNEAL	<u>20MEQ/15ML</u>	<u>A210041</u>	<u>001</u>	Jul 19, 2018
<u>AA</u>			<u>40MEQ/15ML</u>	<u>A210041</u>	<u>002</u>	Jul 19, 2018
<u>AA</u>		ANDA REPOSITORY	<u>20MEQ/15ML</u>	<u>A214892</u>	<u>001</u>	Dec 30, 2022
<u>AA</u>			<u>40MEQ/15ML</u>	<u>A214892</u>	<u>002</u>	Dec 30, 2022
<u>AA</u>		APOTEX	<u>20MEQ/15ML</u>	<u>A211067</u>	<u>001</u>	Aug 08, 2018
<u>AA</u>			<u>40MEQ/15ML</u>	<u>A211067</u>	<u>002</u>	Aug 08, 2018
<u>AA</u>		BELCHER	<u>20MEQ/15ML</u>	<u>A216156</u>	<u>001</u>	Mar 07, 2023
<u>AA</u>			<u>40MEQ/15ML</u>	<u>A216156</u>	<u>002</u>	Mar 07, 2023
<u>AA</u>	+	GENUS LIFESCIENCES	<u>20MEQ/15ML</u>	<u>N206814</u>	<u>001</u>	Dec 22, 2014
<u>AA</u>	+		<u>40MEQ/15ML</u>	<u>N206814</u>	<u>002</u>	Dec 22, 2014
<u>AA</u>		GRANULES	<u>20MEQ/15ML</u>	<u>A213392</u>	<u>001</u>	Jan 29, 2021
<u>AA</u>			<u>40MEQ/15ML</u>	<u>A213392</u>	<u>002</u>	Jan 29, 2021
<u>AA</u>		NOVEL LABS INC	<u>20MEQ/15ML</u>	<u>A209786</u>	<u>001</u>	Aug 29, 2018
<u>AA</u>			<u>40MEQ/15ML</u>	<u>A209786</u>	<u>002</u>	Aug 29, 2018
<u>AA</u>		PHARM ASSOC	<u>20MEQ/15ML</u>	<u>A210766</u>	<u>001</u>	Mar 29, 2019
<u>AA</u>			<u>40MEQ/15ML</u>	<u>A210766</u>	<u>002</u>	Mar 29, 2019
<u>AA</u>		RUBICON	<u>20MEQ/15ML</u>	<u>A214656</u>	<u>001</u>	Jan 13, 2022
<u>AA</u>			<u>40MEQ/15ML</u>	<u>A214656</u>	<u>002</u>	Jan 13, 2022
<u>AA</u>		SCIEGEN PHARMS INC	<u>20MEQ/15ML</u>	<u>A211648</u>	<u>001</u>	May 21, 2021
<u>AA</u>			<u>40MEQ/15ML</u>	<u>A211648</u>	<u>002</u>	May 21, 2021
<u>AA</u>		STRIDES PHARMA	<u>20MEQ/15ML</u>	<u>A211665</u>	<u>002</u>	Jan 16, 2024
<u>AA</u>		WES PHARMA INC	<u>20MEQ/15ML</u>	<u>A213062</u>	<u>001</u>	Jun 06, 2022
<u>AA</u>			<u>40MEQ/15ML</u>	<u>A213062</u>	<u>002</u>	Jun 06, 2022

TABLET, EXTENDED RELEASE; ORAL

KLOR-CON M10

<u>AB1</u>		UPSHER SMITH LABS	<u>10MEQ</u>	<u>A074726</u>	<u>002</u>	Aug 09, 2000
------------	--	-------------------	--------------	----------------	------------	--------------

KLOR-CON M15

<u>AB1</u>		UPSHER SMITH LABS	<u>15MEQ</u>	<u>A074726</u>	<u>003</u>	Jun 06, 2003
------------	--	-------------------	--------------	----------------	------------	--------------

KLOR-CON M20

<u>AB1</u>	!	UPSHER SMITH LABS	<u>20MEQ</u>	<u>A074726</u>	<u>001</u>	Nov 20, 1998
------------	---	-------------------	--------------	----------------	------------	--------------

POTASSIUM CHLORIDE

<u>AB1</u>		ACTAVIS LABS FL INC	<u>10MEQ</u>	<u>A075604</u>	<u>001</u>	Apr 10, 2002
<u>AB1</u>			<u>20MEQ</u>	<u>A075604</u>	<u>002</u>	Apr 10, 2002
<u>AB1</u>		ADARE PHARMS INC	<u>10MEQ</u>	<u>A076368</u>	<u>002</u>	Jun 05, 2019
<u>AB1</u>			<u>20MEQ</u>	<u>A076368</u>	<u>001</u>	Aug 18, 2004
<u>AB1</u>		AMNEAL	<u>10MEQ</u>	<u>A212861</u>	<u>001</u>	May 08, 2020
<u>AB1</u>			<u>20MEQ</u>	<u>A212861</u>	<u>003</u>	May 08, 2020
<u>AB1</u>		ASCENT PHARMS INC	<u>10MEQ</u>	<u>A214422</u>	<u>001</u>	Dec 29, 2020
<u>AB1</u>			<u>15MEQ</u>	<u>A214422</u>	<u>002</u>	Dec 29, 2020
<u>AB1</u>			<u>20MEQ</u>	<u>A214422</u>	<u>003</u>	Dec 29, 2020
<u>AB1</u>		GLENMARK PHARMS LTD	<u>10MEQ</u>	<u>A203562</u>	<u>001</u>	Jul 26, 2016
<u>AB1</u>			<u>20MEQ</u>	<u>A203562</u>	<u>002</u>	Jul 26, 2016

PRESCRIPTION DRUG PRODUCT LIST

POTASSIUM CHLORIDE

TABLET, EXTENDED RELEASE;ORAL

POTASSIUM CHLORIDE

AB1	GRANULES	<u>10MEQ</u>	<u>A214452</u>	<u>001</u>	Oct 21, 2020
AB1		<u>20MEQ</u>	<u>A214452</u>	<u>002</u>	Oct 21, 2020
AB1	GUANGZHOU NOVAKEN	<u>10MEQ</u>	<u>A214395</u>	<u>001</u>	Jan 28, 2021
AB1		<u>20MEQ</u>	<u>A214395</u>	<u>002</u>	Jan 28, 2021
AB1	MACLEODS PHARMS LTD	<u>10MEQ</u>	<u>A216269</u>	<u>001</u>	May 09, 2024
AB1		<u>15MEQ</u>	<u>A216269</u>	<u>002</u>	May 09, 2024
AB1		<u>20MEQ</u>	<u>A216269</u>	<u>003</u>	May 09, 2024
AB1	NOVEL LABS INC	<u>10MEQ</u>	<u>A206347</u>	<u>001</u>	Jan 21, 2016
AB1		<u>20MEQ</u>	<u>A206347</u>	<u>002</u>	Jan 21, 2016
AB1	PRINSTON INC	<u>10MEQ</u>	<u>A209922</u>	<u>001</u>	Apr 30, 2019
AB1		<u>15MEQ</u>	<u>A209922</u>	<u>002</u>	Apr 30, 2019
AB1		<u>20MEQ</u>	<u>A209922</u>	<u>003</u>	Apr 30, 2019
AB1	RUBICON	<u>10MEQ</u>	<u>A216321</u>	<u>001</u>	Jun 16, 2023
AB1		<u>15MEQ</u>	<u>A216321</u>	<u>002</u>	Jun 16, 2023
AB1		<u>20MEQ</u>	<u>A216321</u>	<u>003</u>	Jun 16, 2023
AB1	ZYDUS PHARMS	<u>10MEQ</u>	<u>A210395</u>	<u>001</u>	Sep 17, 2020
AB1		<u>20MEQ</u>	<u>A210395</u>	<u>002</u>	Sep 17, 2020

KLOR-CON

AB2	+ UPSHER SMITH LABS	<u>8MEQ</u>	<u>N019123</u>	<u>001</u>	Apr 17, 1986
AB2	+!	<u>10MEQ</u>	<u>N019123</u>	<u>002</u>	Apr 17, 1986

POTASSIUM CHLORIDE

AB2	AUROBINDO PHARMA	<u>8MEQ</u>	<u>A210921</u>	<u>001</u>	Dec 19, 2018
AB2		<u>10MEQ</u>	<u>A210921</u>	<u>002</u>	Dec 19, 2018
AB2	GRANULES	<u>8MEQ</u>	<u>A211797</u>	<u>001</u>	Mar 04, 2020
AB2		<u>10MEQ</u>	<u>A211797</u>	<u>002</u>	Mar 04, 2020
AB2	MACLEODS PHARMS LTD	<u>8MEQ</u>	<u>A212987</u>	<u>001</u>	Sep 06, 2024
AB2		<u>10MEQ</u>	<u>A212987</u>	<u>002</u>	Sep 06, 2024
AB2	MYLAN	<u>8MEQ</u>	<u>A204662</u>	<u>001</u>	Aug 21, 2014
AB2		<u>10MEQ</u>	<u>A204662</u>	<u>002</u>	Aug 21, 2014
AB2	NOVEL LABS INC	<u>8MEQ</u>	<u>A206759</u>	<u>001</u>	Aug 09, 2016
AB2		<u>10MEQ</u>	<u>A206759</u>	<u>002</u>	Aug 09, 2016
AB2	PADAGIS US	<u>8MEQ</u>	<u>A205993</u>	<u>001</u>	Nov 05, 2015
AB2		<u>10MEQ</u>	<u>A205993</u>	<u>002</u>	Nov 05, 2015
AB2	RISING	<u>8MEQ</u>	<u>A217412</u>	<u>001</u>	Dec 19, 2023
AB2		<u>10MEQ</u>	<u>A217412</u>	<u>002</u>	Dec 19, 2023
AB2	STRIDES PHARMA	<u>8MEQ</u>	<u>A210733</u>	<u>001</u>	Aug 31, 2018
AB2		<u>10MEQ</u>	<u>A210733</u>	<u>002</u>	Aug 31, 2018
AB2	TWI PHARMS	<u>8MEQ</u>	<u>A218979</u>	<u>001</u>	Oct 28, 2024
AB2		<u>10MEQ</u>	<u>A218979</u>	<u>002</u>	Oct 28, 2024
AB2	YICHANG HUMANWELL	<u>8MEQ</u>	<u>A209314</u>	<u>001</u>	Jun 22, 2018
AB2		<u>10MEQ</u>	<u>A209314</u>	<u>002</u>	Jun 22, 2018
AB3	RUBICON	<u>10MEQ</u>	<u>A215725</u>	<u>001</u>	Jul 25, 2022
AB3		<u>20MEQ</u>	<u>A215725</u>	<u>002</u>	Jul 25, 2022
AB3	TWI PHARMS	<u>10MEQ</u>	<u>A209688</u>	<u>001</u>	Jan 12, 2018
AB3	!	<u>20MEQ</u>	<u>A209688</u>	<u>002</u>	Jan 12, 2018
AB3	YICHANG HUMANWELL	<u>10MEQ</u>	<u>A212561</u>	<u>001</u>	Sep 30, 2019
AB3		<u>20MEQ</u>	<u>A212561</u>	<u>002</u>	Sep 30, 2019
	!	RUBICON	15MEQ	A215725	003 Aug 22, 2024

POTASSIUM CHLORIDE; SODIUM CHLORIDE

INJECTABLE; INJECTION

POTASSIUM CHLORIDE 0.149% IN SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER

AP	ICU MEDICAL INC	<u>149MG/100ML; 450MG/100ML</u>	<u>A078446</u>	<u>001</u>	Sep 10, 2008
AP	+! BAXTER HLTHCARE	<u>150MG/100ML; 450MG/100ML</u>	<u>N017648</u>	<u>005</u>	Nov 26, 2002
AP	FRESENIUS KABI USA	<u>150MG/100ML; 450MG/100ML</u>	<u>A212347</u>	<u>001</u>	Sep 17, 2020
AP	+! BAXTER HLTHCARE	<u>150MG/100ML; 900MG/100ML</u>	<u>N017648</u>	<u>001</u>	
AP	FRESENIUS KABI USA	<u>150MG/100ML; 900MG/100ML</u>	<u>A212347</u>	<u>003</u>	Jun 02, 2021
AP	B BRAUN	<u>150MG/100ML; 900MG/100ML</u>	<u>N019708</u>	<u>004</u>	Sep 29, 1989
AP	+! BAXTER HLTHCARE	<u>300MG/100ML; 900MG/100ML</u>	<u>N017648</u>	<u>002</u>	
AP	FRESENIUS KABI USA	<u>300MG/100ML; 900MG/100ML</u>	<u>A212347</u>	<u>002</u>	Sep 17, 2020
AP	ICU MEDICAL INC	<u>149MG/100ML; 900MG/100ML</u>	<u>N019686</u>	<u>001</u>	Oct 17, 1988
AP	ICU MEDICAL INC	<u>298MG/100ML; 900MG/100ML</u>	<u>N019686</u>	<u>002</u>	Oct 17, 1988

PRESCRIPTION DRUG PRODUCT LIST

POTASSIUM CITRATE

TABLET, EXTENDED RELEASE;ORAL

POTASSIUM CITRATE

<u>AB</u>	ANI PHARMS	<u>10MEO</u>	<u>A212779 001</u>	Jan 14, 2020
<u>AB</u>		<u>15MEO</u>	<u>A212779 002</u>	Jan 14, 2020
<u>AB</u>	ASCENT PHARMS INC	<u>5MEO</u>	<u>A214420 001</u>	Feb 05, 2021
<u>AB</u>		<u>10MEO</u>	<u>A214420 002</u>	Feb 05, 2021
<u>AB</u>		<u>15MEO</u>	<u>A214420 003</u>	Feb 05, 2021
<u>AB</u>	BIONPHARMA	<u>10MEO</u>	<u>A212799 001</u>	Jun 29, 2020
<u>AB</u>		<u>15MEO</u>	<u>A212799 002</u>	Jun 29, 2020
<u>AB</u>	HIBROW HLTHCARE	<u>5MEO</u>	<u>A214426 001</u>	Feb 19, 2021
<u>AB</u>		<u>10MEO</u>	<u>A214426 002</u>	Feb 19, 2021
<u>AB</u>		<u>15MEO</u>	<u>A214426 003</u>	Feb 19, 2021
<u>AB</u>	RISING	<u>5MEO</u>	<u>A077440 001</u>	Jun 09, 2006
<u>AB</u>		<u>10MEO</u>	<u>A077440 002</u>	Jun 09, 2006
<u>AB</u>	STRIDES PHARMA	<u>5MEO</u>	<u>A206813 001</u>	Sep 11, 2017
<u>AB</u>		<u>10MEO</u>	<u>A206813 002</u>	Sep 11, 2017
<u>AB</u>		<u>15MEO</u>	<u>A206813 003</u>	Sep 11, 2017
<u>AB</u>	TEVA PHARMS USA INC	<u>5MEO</u>	<u>A209758 001</u>	Mar 05, 2018
<u>AB</u>		<u>10MEO</u>	<u>A209758 002</u>	Mar 05, 2018
<u>AB</u>		<u>15MEO</u>	<u>A209758 003</u>	Mar 05, 2018
<u>AB</u>	ZYDUS PHARMS	<u>5MEO</u>	<u>A203546 001</u>	Aug 06, 2014
<u>AB</u>		<u>10MEO</u>	<u>A203546 002</u>	Aug 06, 2014
<u>AB</u>		<u>15MEO</u>	<u>A203546 003</u>	Aug 06, 2014

UROCIIT-K

<u>AB</u>	+ MISSION PHARMA	<u>5MEO</u>	<u>N019071 001</u>	Aug 30, 1985
<u>AB</u>	+	<u>10MEO</u>	<u>N019071 002</u>	Aug 31, 1992
<u>AB</u>	+!	<u>15MEO</u>	<u>N019071 003</u>	Dec 30, 2009

POTASSIUM PHOSPHATE, DIBASIC; POTASSIUM PHOSPHATE, MONOBASIC

SOLUTION;INTRAVENOUS

POTASSIUM PHOSPHATES

<u>AP</u>	AM REGENT	<u>1.18GM/5ML (236MG/ML) ;1.12GM/5ML (224MG/ML)</u>	<u>A216274 001</u>	Oct 02, 2023
<u>AP</u>		<u>3.54GM/15ML (236MG/ML) ;3.36GM/15ML (224MG/ML)</u>	<u>A216274 002</u>	Oct 02, 2023
<u>AP</u>		<u>11.8GM/50ML (236MG/ML) ;11.2GM/50ML (224MG/ML)</u>	<u>A216274 003</u>	Oct 02, 2023
<u>AP</u>	AMNEAL	<u>1.18GM/5ML (236MG/ML) ;1.12GM/5ML (224MG/ML)</u>	<u>A216344 001</u>	Oct 10, 2023
<u>AP</u>		<u>3.54GM/15ML (236MG/ML) ;3.36GM/15ML (224MG/ML)</u>	<u>A216344 002</u>	Oct 10, 2023
<u>AP</u>		<u>11.8GM/50ML (236MG/ML) ;11.2GM/50ML (224MG/ML)</u>	<u>A216344 003</u>	Oct 10, 2023
<u>AP</u>	+! FRESENIUS KABI USA	<u>1.18GM/5ML (236MG/ML) ;1.12GM/5ML (224MG/ML)</u>	<u>N212832 001</u>	Nov 26, 2019
<u>AP</u>	+!	<u>3.54GM/15ML (236MG/ML) ;3.36GM/15ML (224MG/ML)</u>	<u>N212832 002</u>	Nov 26, 2019
<u>AP</u>	+!	<u>11.8GM/50ML (236MG/ML) ;11.2GM/50ML (224MG/ML)</u>	<u>N212832 003</u>	Nov 26, 2019
<u>AP</u>	SOMERSET THERAPS LLC	<u>1.18GM/5ML (236MG/ML) ;1.12GM/5ML (224MG/ML)</u>	<u>A217726 001</u>	Jul 11, 2024
<u>AP</u>		<u>3.54GM/15ML (236MG/ML) ;3.36GM/15ML (224MG/ML)</u>	<u>A217726 002</u>	Jul 11, 2024
	POTASSIUM PHOSPHATES IN 0.9% SODIUM CHLORIDE			
	+! AMNEAL	<u>1.18GM/250ML (4.72MG/ML) ;1.12GM/250ML (4.48MG/ML)</u>	<u>N218343 001</u>	Jul 26, 2024

POVIDONE-IODINE

SOLUTION/DROPS;OPHTHALMIC

BETADINE

+! ALCON PHARMS LTD

5%

N018634 001 Dec 17, 1986PRALATREXATE

SOLUTION;INTRAVENOUS

FOLOTYN

+ ACROTECH BIOPHARMA

20MG/ML (20MG/ML)

N022468 001 Sep 24, 2009

+!

40MG/2ML (20MG/ML)

N022468 002 Sep 24, 2009PRALIDOXIME CHLORIDE

INJECTABLE;INJECTION

PROTOPAM CHLORIDE

+! BAXTER HLTHCARE

1GM/VIAL

N014134 001

CORP

PRESCRIPTION DRUG PRODUCT LIST

PRALSETINIB

CAPSULE; ORAL

GAVRETO

+! RIGEL PHARMS

100MG

N213721 001 Sep 04, 2020

PRAMIPEXOLE DIHYDROCHLORIDE

TABLET; ORAL

PRAMIPEXOLE DIHYDROCHLORIDE

<u>AB</u>	AUROBINDO PHARMA LTD	<u>0.125MG</u>	<u>A202633 001</u>	Oct 26, 2012
<u>AB</u>		<u>0.25MG</u>	<u>A202633 002</u>	Oct 26, 2012
<u>AB</u>		<u>0.5MG</u>	<u>A202633 003</u>	Oct 26, 2012
<u>AB</u>		<u>0.75MG</u>	<u>A202633 004</u>	Oct 26, 2012
<u>AB</u>		<u>1MG</u>	<u>A202633 005</u>	Oct 26, 2012
<u>AB</u>		<u>1.5MG</u>	<u>A202633 006</u>	Oct 26, 2012
<u>AB</u>	GLENMARK PHARMS LTD	<u>0.125MG</u>	<u>A090781 001</u>	Oct 08, 2010
<u>AB</u>	!	<u>0.25MG</u>	<u>A090781 002</u>	Oct 08, 2010
<u>AB</u>		<u>0.5MG</u>	<u>A090781 003</u>	Oct 08, 2010
<u>AB</u>		<u>0.75MG</u>	<u>A090781 006</u>	Sep 11, 2015
<u>AB</u>		<u>1MG</u>	<u>A090781 004</u>	Oct 08, 2010
<u>AB</u>		<u>1.5MG</u>	<u>A090781 005</u>	Oct 08, 2010
<u>AB</u>	RISING	<u>0.25MG</u>	<u>A211088 001</u>	Oct 03, 2018
<u>AB</u>		<u>0.5MG</u>	<u>A211088 002</u>	Oct 03, 2018
<u>AB</u>		<u>0.75MG</u>	<u>A211088 003</u>	Oct 03, 2018
<u>AB</u>		<u>1MG</u>	<u>A211088 004</u>	Oct 03, 2018
<u>AB</u>		<u>1.5MG</u>	<u>A211088 005</u>	Oct 03, 2018
<u>AB</u>	SCIEGEN PHARMS INC	<u>0.125MG</u>	<u>A203855 001</u>	Oct 28, 2014
<u>AB</u>		<u>0.25MG</u>	<u>A203855 002</u>	Oct 28, 2014
<u>AB</u>		<u>0.5MG</u>	<u>A203855 003</u>	Oct 28, 2014
<u>AB</u>		<u>0.75MG</u>	<u>A203855 004</u>	Oct 28, 2014
<u>AB</u>		<u>1MG</u>	<u>A203855 005</u>	Oct 28, 2014
<u>AB</u>		<u>1.5MG</u>	<u>A203855 006</u>	Oct 28, 2014
<u>AB</u>	STRIDES PHARMA	<u>0.125MG</u>	<u>A202702 001</u>	Jun 03, 2014
<u>AB</u>		<u>0.25MG</u>	<u>A202702 002</u>	Jun 03, 2014
<u>AB</u>		<u>0.5MG</u>	<u>A202702 003</u>	Jun 03, 2014
<u>AB</u>		<u>0.75MG</u>	<u>A202702 004</u>	Jun 03, 2014
<u>AB</u>		<u>1MG</u>	<u>A202702 005</u>	Jun 03, 2014
<u>AB</u>		<u>1.5MG</u>	<u>A202702 006</u>	Jun 03, 2014
<u>AB</u>	TORRENT PHARMS	<u>0.125MG</u>	<u>A090865 001</u>	Oct 08, 2010
<u>AB</u>		<u>0.25MG</u>	<u>A090865 002</u>	Oct 08, 2010
<u>AB</u>		<u>0.5MG</u>	<u>A090865 003</u>	Oct 08, 2010
<u>AB</u>		<u>0.75MG</u>	<u>A090865 004</u>	Oct 08, 2010
<u>AB</u>		<u>1MG</u>	<u>A090865 005</u>	Oct 08, 2010
<u>AB</u>		<u>1.5MG</u>	<u>A090865 006</u>	Oct 08, 2010
<u>AB</u>	ZENNOVA	<u>0.125MG</u>	<u>A090151 001</u>	Apr 30, 2012
<u>AB</u>		<u>0.25MG</u>	<u>A090151 002</u>	Apr 30, 2012
<u>AB</u>		<u>0.5MG</u>	<u>A090151 003</u>	Apr 30, 2012
<u>AB</u>		<u>0.75MG</u>	<u>A090151 006</u>	Apr 30, 2012
<u>AB</u>		<u>1MG</u>	<u>A090151 004</u>	Apr 30, 2012
<u>AB</u>		<u>1.5MG</u>	<u>A090151 005</u>	Apr 30, 2012
<u>AB</u>	ZYDUS PHARMS USA INC	<u>0.125MG</u>	<u>A078920 001</u>	Jul 06, 2010
<u>AB</u>		<u>0.25MG</u>	<u>A078920 002</u>	Jul 06, 2010
<u>AB</u>		<u>0.5MG</u>	<u>A078920 003</u>	Jul 06, 2010
<u>AB</u>		<u>0.75MG</u>	<u>A078920 006</u>	Nov 23, 2022
<u>AB</u>		<u>1MG</u>	<u>A078920 004</u>	Jul 06, 2010
<u>AB</u>		<u>1.5MG</u>	<u>A078920 005</u>	Jul 06, 2010

TABLET, EXTENDED RELEASE; ORAL

PRAMIPEXOLE DIHYDROCHLORIDE

<u>AB</u>	ACTAVIS ELIZABETH	<u>0.375MG</u>	<u>A201963 001</u>	Apr 21, 2016
<u>AB</u>		<u>0.75MG</u>	<u>A201963 002</u>	Apr 21, 2016
<u>AB</u>		<u>1.5MG</u>	<u>A201963 003</u>	Apr 21, 2016
<u>AB</u>		<u>2.25MG</u>	<u>A203615 001</u>	Oct 14, 2016
<u>AB</u>		<u>3MG</u>	<u>A201963 004</u>	Apr 21, 2016
<u>AB</u>		<u>3.75MG</u>	<u>A203615 002</u>	Jan 03, 2017
<u>AB</u>		<u>4.5MG</u>	<u>A201963 005</u>	Apr 21, 2016
<u>AB</u>	ALEMBIC	<u>0.375MG</u>	<u>A204518 001</u>	Jan 02, 2019
<u>AB</u>		<u>0.75MG</u>	<u>A204518 002</u>	Jan 02, 2019
<u>AB</u>		<u>1.5MG</u>	<u>A204518 003</u>	Jan 02, 2019
<u>AB</u>		<u>2.25MG</u>	<u>A204518 004</u>	Jan 02, 2019
<u>AB</u>		<u>3MG</u>	<u>A204518 005</u>	Jan 02, 2019
<u>AB</u>		<u>3.75MG</u>	<u>A204518 006</u>	Jan 02, 2019
<u>AB</u>		<u>4.5MG</u>	<u>A204518 007</u>	Jan 02, 2019

PRESCRIPTION DRUG PRODUCT LIST

PRAMIPEXOLE DIHYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL

PRAMIPEXOLE DIHYDROCHLORIDE

<u>AB</u>	CHARTWELL RX	<u>0.375MG</u>	<u>A202353 001</u>	Dec 04, 2014
<u>AB</u>		<u>0.75MG</u>	<u>A202353 002</u>	Dec 04, 2014
<u>AB</u>		<u>1.5MG</u>	<u>A202353 003</u>	Dec 04, 2014
<u>AB</u>		<u>3MG</u>	<u>A202353 004</u>	Dec 04, 2014
<u>AB</u>		<u>4.5MG</u>	<u>A202353 005</u>	Dec 04, 2014
<u>AB</u>	! DR REDDYS	<u>0.375MG</u>	<u>A203354 001</u>	Aug 07, 2015
<u>AB</u>		<u>0.75MG</u>	<u>A203354 002</u>	Aug 07, 2015
<u>AB</u>		<u>1.5MG</u>	<u>A203354 003</u>	Aug 07, 2015
<u>AB</u>		<u>3MG</u>	<u>A203354 004</u>	Aug 07, 2015
<u>AB</u>		<u>4.5MG</u>	<u>A203354 005</u>	Aug 07, 2015
<u>AB</u>	MACLEODS PHARMS LTD	<u>0.375MG</u>	<u>A206156 001</u>	Jun 24, 2016
<u>AB</u>		<u>0.75MG</u>	<u>A206156 002</u>	Jun 24, 2016
<u>AB</u>		<u>1.5MG</u>	<u>A206156 003</u>	Jun 24, 2016
<u>AB</u>		<u>2.25MG</u>	<u>A206156 004</u>	Jun 24, 2016
<u>AB</u>		<u>3MG</u>	<u>A206156 005</u>	Jun 24, 2016
<u>AB</u>		<u>3.75MG</u>	<u>A206156 007</u>	Jan 23, 2017
<u>AB</u>		<u>4.5MG</u>	<u>A206156 006</u>	Jun 24, 2016
<u>AB</u>	NOVAST LABS	<u>0.375MG</u>	<u>A213444 001</u>	Feb 03, 2022
<u>AB</u>		<u>0.75MG</u>	<u>A213444 002</u>	Feb 03, 2022
<u>AB</u>		<u>1.5MG</u>	<u>A213444 003</u>	Feb 03, 2022
<u>AB</u>		<u>2.25MG</u>	<u>A213444 004</u>	Feb 03, 2022
<u>AB</u>		<u>3MG</u>	<u>A213444 005</u>	Feb 03, 2022
<u>AB</u>		<u>3.75MG</u>	<u>A213444 006</u>	Feb 03, 2022
<u>AB</u>		<u>4.5MG</u>	<u>A213444 007</u>	Feb 03, 2022
<u>AB</u>	XIAMEN LP PHARM CO	<u>0.375MG</u>	<u>A212797 001</u>	Jun 11, 2021
<u>AB</u>		<u>0.75MG</u>	<u>A212797 002</u>	Jun 11, 2021
<u>AB</u>	ZYDUS PHARMS	<u>0.375MG</u>	<u>A202891 001</u>	Dec 12, 2017
<u>AB</u>		<u>0.75MG</u>	<u>A202891 002</u>	Dec 12, 2017
<u>AB</u>		<u>1.5MG</u>	<u>A202891 003</u>	Dec 12, 2017
<u>AB</u>		<u>2.25MG</u>	<u>A202891 004</u>	Dec 12, 2017
<u>AB</u>		<u>3MG</u>	<u>A202891 005</u>	Dec 12, 2017
<u>AB</u>		<u>3.75MG</u>	<u>A202891 006</u>	Dec 12, 2017
<u>AB</u>		<u>4.5MG</u>	<u>A202891 007</u>	Dec 12, 2017

PRAMLINTIDE ACETATE

INJECTABLE;SUBCUTANEOUS

SYMLIN

+	ASTRAZENECA AB	EQ 1.5MG BASE/1.5ML (EQ 1MG BASE/ML)	N021332 002	Sep 25, 2007
+	!	EQ 2.7MG BASE/2.7ML (EQ 1MG BASE/ML)	N021332 003	Sep 25, 2007

PRASTERONE

INSERT;VAGINAL

INTRAROSA

+	!	MILLICENT	6.5MG	N208470 001	Nov 16, 2016
---	---	-----------	-------	-------------	--------------

PRASUGREL HYDROCHLORIDE

TABLET;ORAL

EFFIENT

<u>AB</u>	+	COSETTE	<u>EQ 5MG BASE</u>	<u>N022307 001</u>	Jul 10, 2009
<u>AB</u>	+	!	<u>EQ 10MG BASE</u>	<u>N022307 002</u>	Jul 10, 2009

PRASUGREL

<u>AB</u>	ACCORD HLTHCARE	<u>EQ 5MG BASE</u>	<u>A205987 001</u>	Feb 02, 2018
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A205987 002</u>	Feb 02, 2018
<u>AB</u>	AMNEAL PHARMS	<u>EQ 5MG BASE</u>	<u>A205913 001</u>	Jun 19, 2018
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A205913 002</u>	Jun 19, 2018
<u>AB</u>	AUROBINDO PHARMA	<u>EQ 5MG BASE</u>	<u>A205888 001</u>	Oct 16, 2017
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A205888 002</u>	Oct 16, 2017
<u>AB</u>	HEC PHARM	<u>EQ 5MG BASE</u>	<u>A206021 001</u>	Jan 16, 2019
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A206021 002</u>	Jan 16, 2019
<u>AB</u>	MYLAN	<u>EQ 5MG BASE</u>	<u>A205927 001</u>	Jul 12, 2017
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A205927 002</u>	Jul 12, 2017
<u>AB</u>	PANACEA	<u>EQ 5MG BASE</u>	<u>A205897 001</u>	Oct 16, 2017
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A205897 002</u>	Oct 16, 2017

PRASUGREL HYDROCHLORIDE

<u>AB</u>	UNICHEM	<u>EQ 5MG BASE</u>	<u>A213315 001</u>	Aug 28, 2023
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A213315 002</u>	Aug 28, 2023

PRESCRIPTION DRUG PRODUCT LIST

PRAVASTATIN SODIUM

TABLET; ORAL

PRAVASTATIN SODIUM

<u>AB</u>	ACCORD HLTHCARE	<u>10MG</u>	<u>A207068 001</u>	Nov 17, 2016
<u>AB</u>		<u>20MG</u>	<u>A207068 002</u>	Nov 17, 2016
<u>AB</u>		<u>40MG</u>	<u>A207068 003</u>	Nov 17, 2016
<u>AB</u>		<u>80MG</u>	<u>A207068 004</u>	Nov 17, 2016
<u>AB</u>	APNAR PHARMA LP	<u>10MG</u>	<u>A077491 002</u>	Oct 23, 2006
<u>AB</u>		<u>20MG</u>	<u>A077491 003</u>	Oct 23, 2006
<u>AB</u>		<u>40MG</u>	<u>A077491 004</u>	Oct 23, 2006
<u>AB</u>		<u>80MG</u>	<u>A077491 001</u>	Feb 11, 2008
<u>AB</u>	APOTEX	<u>10MG</u>	<u>A076341 001</u>	Oct 23, 2006
<u>AB</u>		<u>20MG</u>	<u>A076341 002</u>	Oct 23, 2006
<u>AB</u>		<u>40MG</u>	<u>A076341 003</u>	Oct 23, 2006
<u>AB</u>		<u>80MG</u>	<u>A076341 004</u>	Dec 28, 2007
<u>AB</u>	AUROBINDO PHARMA	<u>10MG</u>	<u>A203367 001</u>	Feb 02, 2017
<u>AB</u>		<u>20MG</u>	<u>A203367 002</u>	Feb 02, 2017
<u>AB</u>		<u>40MG</u>	<u>A203367 003</u>	Feb 02, 2017
<u>AB</u>		<u>80MG</u>	<u>A203367 004</u>	Feb 02, 2017
<u>AB</u>	AUROBINDO PHARMA USA	<u>10MG</u>	<u>A076056 001</u>	Apr 24, 2006
<u>AB</u>		<u>20MG</u>	<u>A076056 002</u>	Apr 24, 2006
<u>AB</u>		<u>40MG</u>	<u>A076056 003</u>	Apr 24, 2006
<u>AB</u>	!	<u>80MG</u>	<u>A077793 001</u>	Jan 15, 2008
<u>AB</u>	BIOCON PHARMA	<u>10MG</u>	<u>A209869 001</u>	Apr 13, 2018
<u>AB</u>		<u>20MG</u>	<u>A209869 002</u>	Apr 13, 2018
<u>AB</u>		<u>40MG</u>	<u>A209869 003</u>	Apr 13, 2018
<u>AB</u>		<u>80MG</u>	<u>A209869 004</u>	Apr 13, 2018
<u>AB</u>	CHARTWELL RX	<u>10MG</u>	<u>A077917 001</u>	Jan 08, 2008
<u>AB</u>		<u>20MG</u>	<u>A077917 002</u>	Jan 08, 2008
<u>AB</u>		<u>40MG</u>	<u>A077917 003</u>	Jan 08, 2008
<u>AB</u>		<u>80MG</u>	<u>A077917 004</u>	Jan 08, 2008
<u>AB</u>	CIPLA	<u>10MG</u>	<u>A077904 001</u>	Oct 23, 2006
<u>AB</u>		<u>20MG</u>	<u>A077904 002</u>	Oct 23, 2006
<u>AB</u>		<u>40MG</u>	<u>A077904 003</u>	Oct 23, 2006
<u>AB</u>		<u>80MG</u>	<u>A077904 004</u>	Mar 22, 2016
<u>AB</u>	DR REDDYS LABS INC	<u>10MG</u>	<u>A076714 001</u>	Oct 23, 2006
<u>AB</u>		<u>20MG</u>	<u>A076714 002</u>	Oct 23, 2006
<u>AB</u>		<u>40MG</u>	<u>A076714 003</u>	Oct 23, 2006
<u>AB</u>		<u>80MG</u>	<u>A076714 004</u>	Dec 28, 2007
<u>AB</u>	GLENMARK PHARMS LTD	<u>10MG</u>	<u>A077987 001</u>	May 11, 2007
<u>AB</u>		<u>20MG</u>	<u>A077987 002</u>	May 11, 2007
<u>AB</u>		<u>40MG</u>	<u>A077987 003</u>	May 11, 2007
<u>AB</u>		<u>80MG</u>	<u>A077987 004</u>	Dec 28, 2007
<u>AB</u>	WATSON LABS	<u>10MG</u>	<u>A076939 004</u>	Oct 23, 2006
<u>AB</u>		<u>20MG</u>	<u>A076939 003</u>	Oct 23, 2006
<u>AB</u>		<u>40MG</u>	<u>A076939 002</u>	Oct 23, 2006
<u>AB</u>		<u>80MG</u>	<u>A076939 001</u>	Dec 28, 2007
<u>AB</u>	ZYDUS PHARMS USA	<u>20MG</u>	<u>A077751 002</u>	Apr 30, 2008
<u>AB</u>		<u>40MG</u>	<u>A077751 003</u>	Apr 30, 2008
<u>AB</u>		<u>80MG</u>	<u>A077751 004</u>	Apr 30, 2008

PRAZIOUANTEL

TABLET; ORAL

PRAZIOUANTEL

! ENDO OPERATIONS

600MG

A208820 001 Nov 27, 2017

PRAZOSIN HYDROCHLORIDE

CAPSULE; ORAL

MINIPRESS

<u>AB</u>	+	PFIZER	<u>EQ 1MG BASE</u>	<u>N017442 002</u>
<u>AB</u>	+	!	<u>EQ 2MG BASE</u>	<u>N017442 003</u>
<u>AB</u>	+		<u>EQ 5MG BASE</u>	<u>N017442 001</u>

PRAZOSIN HYDROCHLORIDE

<u>AB</u>	ALEMBIC	<u>EQ 1MG BASE</u>	<u>A217268 001</u>	Mar 06, 2023
<u>AB</u>		<u>EQ 2MG BASE</u>	<u>A217268 002</u>	Mar 06, 2023
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A217268 003</u>	Mar 06, 2023
<u>AB</u>	APPCO	<u>EQ 1MG BASE</u>	<u>A213406 001</u>	Oct 21, 2022
<u>AB</u>		<u>EQ 2MG BASE</u>	<u>A213406 002</u>	Oct 21, 2022
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A213406 003</u>	Oct 21, 2022
<u>AB</u>	AUROBINDO PHARMA LTD	<u>EQ 1MG BASE</u>	<u>A213052 001</u>	Mar 31, 2023
<u>AB</u>		<u>EQ 2MG BASE</u>	<u>A213052 002</u>	Mar 31, 2023
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A213052 003</u>	Mar 31, 2023

PRESCRIPTION DRUG PRODUCT LIST

PRAZOSIN HYDROCHLORIDE

CAPSULE; ORAL

PRAZOSIN HYDROCHLORIDE

<u>AB</u>		GRANULES	<u>EQ 1MG BASE</u>	<u>A214608 001</u>	Dec 23, 2021
<u>AB</u>			<u>EQ 2MG BASE</u>	<u>A214608 002</u>	Dec 23, 2021
<u>AB</u>			<u>EQ 5MG BASE</u>	<u>A214608 003</u>	Dec 23, 2021
<u>AB</u>		MANKIND PHARMA	<u>EQ 1MG BASE</u>	<u>A215697 001</u>	Dec 30, 2022
<u>AB</u>			<u>EQ 2MG BASE</u>	<u>A215697 002</u>	Dec 30, 2022
<u>AB</u>			<u>EQ 5MG BASE</u>	<u>A215697 003</u>	Dec 30, 2022
<u>AB</u>		MSN	<u>EQ 1MG BASE</u>	<u>A216727 001</u>	Sep 17, 2024
<u>AB</u>			<u>EQ 2MG BASE</u>	<u>A216727 002</u>	Sep 17, 2024
<u>AB</u>			<u>EQ 5MG BASE</u>	<u>A216727 003</u>	Sep 17, 2024
<u>AB</u>		MYLAN	<u>EQ 1MG BASE</u>	<u>A072575 003</u>	May 16, 1989
<u>AB</u>			<u>EQ 2MG BASE</u>	<u>A072575 002</u>	May 16, 1989
<u>AB</u>			<u>EQ 5MG BASE</u>	<u>A072575 001</u>	May 16, 1989
<u>AB</u>		NOVITIUM PHARMA	<u>EQ 1MG BASE</u>	<u>A210971 001</u>	Oct 03, 2018
<u>AB</u>			<u>EQ 2MG BASE</u>	<u>A210971 002</u>	Oct 03, 2018
<u>AB</u>			<u>EQ 5MG BASE</u>	<u>A210971 003</u>	Oct 03, 2018
<u>AB</u>		TEVA PHARMS	<u>EQ 1MG BASE</u>	<u>A071745 002</u>	Sep 12, 1988
<u>AB</u>			<u>EQ 2MG BASE</u>	<u>A071745 003</u>	Sep 12, 1988
<u>AB</u>			<u>EQ 5MG BASE</u>	<u>A071745 001</u>	Sep 12, 1988

PREDNICARBATE

OINTMENT; TOPICAL

PREDNICARBATE

! FOUGERA PHARMS

0.1%

A077236 001 Mar 09, 2007

PREDNISOLONE

SYRUP; ORAL

PREDNISOLONE

<u>AA</u>	!	CHARTWELL RX	<u>15MG/5ML</u>	<u>A040323 001</u>	May 13, 1999
<u>AA</u>		LANNETT CO INC	<u>15MG/5ML</u>	<u>A040775 001</u>	Sep 21, 2007
<u>AA</u>		PHARM ASSOC	<u>15MG/5ML</u>	<u>A040571 001</u>	Aug 25, 2005

PRELONE

<u>AA</u>		TEVA	<u>15MG/5ML</u>	<u>A089081 001</u>	Feb 04, 1986
-----------	--	------	-----------------	--------------------	--------------

TABLET; ORAL

PREDNISOLONE

<u>AB</u>		AUROBINDO PHARMA LTD	<u>5MG</u>	<u>A215673 001</u>	Mar 17, 2023
<u>AB</u>	+	WATSON LABS	<u>5MG</u>	<u>A080354 001</u>	
<u>AB</u>		ZHEJIANG XIANJU	<u>5MG</u>	<u>A218083 001</u>	May 02, 2024
<u>BX</u>		CHARTWELL MOLECULAR	<u>5MG</u>	A080531 002	

PREDNISOLONE ACETATE

SUSPENSION/DROPS; OPHTHALMIC

OMNIPRED

<u>AB</u>	+	SANDOZ	<u>1%</u>	<u>N017469 001</u>	
-----------	---	--------	-----------	--------------------	--

PRED FORTE

<u>AB</u>	+	ABBVIE	<u>1%</u>	<u>N017011 001</u>	
-----------	---	--------	-----------	--------------------	--

PREDNISOLONE ACETATE

<u>AB</u>		LUPIN LTD	<u>1%</u>	<u>A216935 001</u>	Aug 02, 2024
		PRED MILD			
	+	ABBVIE	<u>0.12%</u>	N017100 001	

PREDNISOLONE SODIUM PHOSPHATE

SOLUTION; ORAL

PEDIAAPRED

<u>AA</u>	+	SETON PHARM	<u>EQ 5MG BASE/5ML</u>	<u>N019157 001</u>	May 28, 1986
-----------	---	-------------	------------------------	--------------------	--------------

PREDNISOLONE SODIUM PHOSPHATE

<u>AA</u>		AMNEAL	<u>EQ 15MG BASE/5ML</u>	<u>A216715 001</u>	Oct 25, 2022
<u>AA</u>		CHARTWELL RX	<u>EQ 5MG BASE/5ML</u>	<u>A075988 001</u>	May 25, 2004
<u>AA</u>		EDENBRIDGE PHARMS	<u>EQ 10MG BASE/5ML</u>	<u>A203559 001</u>	Dec 20, 2016
<u>AA</u>			<u>EQ 15MG BASE/5ML</u>	<u>A203559 003</u>	Feb 06, 2023
<u>AA</u>			<u>EQ 20MG BASE/5ML</u>	<u>A203559 002</u>	Dec 20, 2016
<u>AA</u>			<u>EQ 25MG BASE/5ML</u>	<u>A203559 004</u>	Feb 06, 2023
<u>AA</u>	!	MISSION PHARMA	<u>EQ 25MG BASE/5ML</u>	<u>A091396 001</u>	Sep 13, 2010
<u>AA</u>	!	PHARM ASSOC	<u>EQ 10MG BASE/5ML</u>	<u>A078465 001</u>	Mar 07, 2008
<u>AA</u>	!		<u>EQ 15MG BASE/5ML</u>	<u>A076913 001</u>	Apr 25, 2005
<u>AA</u>	!		<u>EQ 20MG BASE/5ML</u>	<u>A078988 001</u>	Jun 09, 2008

SOLUTION/DROPS; OPHTHALMIC

PREDNISOLONE SODIUM PHOSPHATE

! BAUSCH AND LOMB

EQ 0.9% PHOSPHATE

A040070 001 Jul 29, 1994

PRESCRIPTION DRUG PRODUCT LIST

PREDNISOLONE SODIUM PHOSPHATE

TABLET, ORALLY DISINTEGRATING;ORAL
ORAPRED ODT

+	ADVANZ PHARMA	EQ 10MG BASE	N021959 001	Jun 01, 2006
+		EQ 15MG BASE	N021959 002	Jun 01, 2006
+	!	EQ 30MG BASE	N021959 003	Jun 01, 2006

PREDNISOLONE SODIUM PHOSPHATE; SULFACETAMIDE SODIUM

SOLUTION/DROPS;OPHTHALMIC

SULFACETAMIDE SODIUM AND PREDNISOLONE SODIUM PHOSPHATE
! BAUSCH AND LOMB EQ 0.23% PHOSPHATE;10%

A074449 001 Dec 29, 1995

PREDNISON

SOLUTION;ORAL

PREDNISON

+	!	HIKMA	5MG/5ML	A088703 001	Nov 08, 1984
---	---	-------	---------	-------------	--------------

PREDNISON INTENSOL

+	!	HIKMA	5MG/ML	A088810 001	Feb 20, 1985
---	---	-------	--------	-------------	--------------

TABLET;ORAL

PREDNISON

AB		ALLIED	5MG	A212629 001	Dec 05, 2023	
AB			10MG	A212629 002	Dec 05, 2023	
AB			20MG	A212629 003	Dec 05, 2023	
AB		AMNEAL	1MG	A213385 001	Jun 16, 2020	
AB			5MG	A213385 002	Jun 16, 2020	
AB			10MG	A213386 001	Jun 24, 2020	
AB			20MG	A213386 002	Jun 24, 2020	
AB		AUROBINDO PHARMA LTD	1MG	A215671 001	Nov 16, 2021	
AB			2.5MG	A215672 001	Mar 28, 2022	
AB			5MG	A215672 002	Mar 28, 2022	
AB			10MG	A215672 003	Mar 28, 2022	
AB			20MG	A215672 004	Mar 28, 2022	
AB			50MG	A215672 005	Mar 28, 2022	
AB		ENDO OPERATIONS	1MG	A040584 001	Dec 21, 2004	
AB			2.5MG	A040581 001	Dec 21, 2004	
AB			5MG	A040256 001	Jul 12, 2002	
AB			10MG	A040256 002	Jul 12, 2002	
AB			20MG	A040392 001	Feb 12, 2003	
AB	+	!	HIKMA	1MG	A087800 001	Apr 22, 1982
AB	+	!		2.5MG	A087801 001	Apr 22, 1982
AB	+	!		5MG	A080352 001	
AB	+	!		10MG	A084122 001	
AB	+	!		20MG	A087342 001	
AB	+	!		50MG	A084283 001	
AB		JUBILANT CADISTA	1MG	A040611 001	Jun 06, 2005	
AB			2.5MG	A040362 004	Apr 17, 2023	
AB			5MG	A040362 002	Aug 29, 2001	
AB			10MG	A040362 001	Aug 29, 2001	
AB			20MG	A040362 003	Jun 29, 2005	
AB			50MG	A040362 005	Apr 17, 2023	
AB		MYLAN	5MG	A080292 001		
AB			10MG	A088832 001	Dec 04, 1985	
AB			20MG	A083677 001		
AB		NOVITIUM PHARMA	1MG	A215246 001	Jul 06, 2021	
AB			2.5MG	A211575 001	Nov 15, 2019	
AB			5MG	A211575 002	Nov 15, 2019	
AB			10MG	A211575 003	Nov 15, 2019	
AB			20MG	A211575 004	Nov 15, 2019	
AB			50MG	A211575 005	Nov 15, 2019	
AB		STRIDES PHARMA	1MG	A210785 001	Sep 02, 2020	
AB			2.5MG	A208412 004	Nov 20, 2020	
AB			5MG	A208412 005	Nov 20, 2020	
AB			10MG	A208412 001	Feb 11, 2021	
AB			20MG	A208412 002	Feb 11, 2021	
AB			50MG	A208412 003	Jan 11, 2022	
AB		SUN PHARM INDUSTRIES	5MG	A089247 002	Dec 04, 1985	
AB			10MG	A089247 003	Dec 04, 1985	
AB			20MG	A089247 001	Dec 04, 1985	
AB		WATSON LABS	5MG	A080356 001		
AB			10MG	A085162 001		
AB			20MG	A085161 001		
BX		CHARTWELL MOLECULAR	20MG	A084275 001		

PRESCRIPTION DRUG PRODUCT LIST

PREGABALIN

CAPSULE; ORAL

LYRICA

<u>AB</u>	+	UPJOHN	<u>25MG</u>	<u>N021446</u>	<u>001</u>	Dec 30, 2004
<u>AB</u>	+		<u>50MG</u>	<u>N021446</u>	<u>002</u>	Dec 30, 2004
<u>AB</u>	+		<u>75MG</u>	<u>N021446</u>	<u>003</u>	Dec 30, 2004
<u>AB</u>	+		<u>100MG</u>	<u>N021446</u>	<u>004</u>	Dec 30, 2004
<u>AB</u>	+		<u>150MG</u>	<u>N021446</u>	<u>005</u>	Dec 30, 2004
<u>AB</u>	+		<u>200MG</u>	<u>N021446</u>	<u>006</u>	Dec 30, 2004
<u>AB</u>	+		<u>225MG</u>	<u>N021446</u>	<u>007</u>	Dec 30, 2004
<u>AB</u>	+		<u>300MG</u>	<u>N021446</u>	<u>008</u>	Dec 30, 2004

PREGABALIN

<u>AB</u>		ACTAVIS ELIZABETH	<u>25MG</u>	<u>A091025</u>	<u>001</u>	Jul 09, 2020
<u>AB</u>			<u>50MG</u>	<u>A091025</u>	<u>002</u>	Jul 09, 2020
<u>AB</u>			<u>75MG</u>	<u>A091025</u>	<u>003</u>	Jul 09, 2020
<u>AB</u>			<u>100MG</u>	<u>A091025</u>	<u>004</u>	Jul 09, 2020
<u>AB</u>			<u>150MG</u>	<u>A091025</u>	<u>005</u>	Jul 09, 2020
<u>AB</u>			<u>200MG</u>	<u>A091025</u>	<u>006</u>	Jul 09, 2020
<u>AB</u>			<u>225MG</u>	<u>A091025</u>	<u>007</u>	Jul 09, 2020
<u>AB</u>			<u>300MG</u>	<u>A091025</u>	<u>008</u>	Jul 09, 2020
<u>AB</u>		ADAPTIS	<u>25MG</u>	<u>A208113</u>	<u>001</u>	May 17, 2024
<u>AB</u>			<u>25MG</u>	<u>A216197</u>	<u>001</u>	Jul 18, 2022
<u>AB</u>			<u>50MG</u>	<u>A208113</u>	<u>002</u>	May 17, 2024
<u>AB</u>			<u>50MG</u>	<u>A216197</u>	<u>002</u>	Jul 18, 2022
<u>AB</u>			<u>75MG</u>	<u>A208113</u>	<u>004</u>	May 17, 2024
<u>AB</u>			<u>75MG</u>	<u>A216197</u>	<u>003</u>	Jul 18, 2022
<u>AB</u>			<u>100MG</u>	<u>A208113</u>	<u>005</u>	May 17, 2024
<u>AB</u>			<u>100MG</u>	<u>A216197</u>	<u>004</u>	Jul 18, 2022
<u>AB</u>			<u>150MG</u>	<u>A208113</u>	<u>003</u>	May 17, 2024
<u>AB</u>			<u>150MG</u>	<u>A216197</u>	<u>005</u>	Jul 18, 2022
<u>AB</u>			<u>200MG</u>	<u>A208113</u>	<u>006</u>	May 17, 2024
<u>AB</u>			<u>200MG</u>	<u>A216197</u>	<u>006</u>	Jul 18, 2022
<u>AB</u>			<u>225MG</u>	<u>A208113</u>	<u>007</u>	May 17, 2024
<u>AB</u>			<u>225MG</u>	<u>A216197</u>	<u>007</u>	Jul 18, 2022
<u>AB</u>			<u>300MG</u>	<u>A208113</u>	<u>008</u>	May 17, 2024
<u>AB</u>			<u>300MG</u>	<u>A216197</u>	<u>008</u>	Jul 18, 2022
<u>AB</u>		ALEMBIC	<u>25MG</u>	<u>A203459</u>	<u>001</u>	Jul 19, 2019
<u>AB</u>			<u>50MG</u>	<u>A203459</u>	<u>002</u>	Jul 19, 2019
<u>AB</u>			<u>75MG</u>	<u>A203459</u>	<u>003</u>	Jul 19, 2019
<u>AB</u>			<u>100MG</u>	<u>A203459</u>	<u>004</u>	Jul 19, 2019
<u>AB</u>			<u>150MG</u>	<u>A203459</u>	<u>005</u>	Jul 19, 2019
<u>AB</u>			<u>200MG</u>	<u>A203459</u>	<u>006</u>	Jul 19, 2019
<u>AB</u>			<u>225MG</u>	<u>A203459</u>	<u>007</u>	Jul 19, 2019
<u>AB</u>			<u>300MG</u>	<u>A203459</u>	<u>008</u>	Jul 19, 2019
<u>AB</u>		ALKEM LABS LTD	<u>25MG</u>	<u>A207799</u>	<u>007</u>	Sep 30, 2019
<u>AB</u>			<u>50MG</u>	<u>A207799</u>	<u>008</u>	Sep 30, 2019
<u>AB</u>			<u>75MG</u>	<u>A207799</u>	<u>001</u>	Jul 19, 2019
<u>AB</u>			<u>100MG</u>	<u>A207799</u>	<u>002</u>	Jul 19, 2019
<u>AB</u>			<u>150MG</u>	<u>A207799</u>	<u>003</u>	Jul 19, 2019
<u>AB</u>			<u>200MG</u>	<u>A207799</u>	<u>004</u>	Jul 19, 2019
<u>AB</u>			<u>225MG</u>	<u>A207799</u>	<u>005</u>	Jul 19, 2019
<u>AB</u>			<u>300MG</u>	<u>A207799</u>	<u>006</u>	Jul 19, 2019
<u>AB</u>		AMNEAL PHARMS CO	<u>25MG</u>	<u>A209743</u>	<u>001</u>	Jul 19, 2019
<u>AB</u>			<u>50MG</u>	<u>A209743</u>	<u>002</u>	Jul 19, 2019
<u>AB</u>			<u>75MG</u>	<u>A209743</u>	<u>003</u>	Jul 19, 2019
<u>AB</u>			<u>100MG</u>	<u>A209743</u>	<u>004</u>	Jul 19, 2019
<u>AB</u>			<u>150MG</u>	<u>A209743</u>	<u>005</u>	Jul 19, 2019
<u>AB</u>			<u>200MG</u>	<u>A209743</u>	<u>006</u>	Jul 19, 2019
<u>AB</u>			<u>225MG</u>	<u>A209743</u>	<u>007</u>	Jul 19, 2019
<u>AB</u>			<u>300MG</u>	<u>A209743</u>	<u>008</u>	Jul 19, 2019
<u>AB</u>		APOTEX	<u>25MG</u>	<u>A211685</u>	<u>001</u>	Jul 07, 2021
<u>AB</u>			<u>50MG</u>	<u>A211685</u>	<u>002</u>	Jul 07, 2021
<u>AB</u>			<u>75MG</u>	<u>A211685</u>	<u>003</u>	Jul 07, 2021
<u>AB</u>			<u>100MG</u>	<u>A211685</u>	<u>004</u>	Jul 07, 2021
<u>AB</u>			<u>150MG</u>	<u>A211685</u>	<u>005</u>	Jul 07, 2021
<u>AB</u>			<u>200MG</u>	<u>A211685</u>	<u>006</u>	Jul 07, 2021
<u>AB</u>			<u>225MG</u>	<u>A211685</u>	<u>007</u>	Jul 07, 2021
<u>AB</u>			<u>300MG</u>	<u>A211685</u>	<u>008</u>	Jul 07, 2021
<u>AB</u>		AUROBINDO PHARMA	<u>25MG</u>	<u>A205321</u>	<u>001</u>	Mar 29, 2023
<u>AB</u>			<u>50MG</u>	<u>A205321</u>	<u>002</u>	Mar 29, 2023
<u>AB</u>			<u>75MG</u>	<u>A205321</u>	<u>003</u>	Mar 29, 2023
<u>AB</u>			<u>100MG</u>	<u>A205321</u>	<u>004</u>	Mar 29, 2023

PRESCRIPTION DRUG PRODUCT LIST

PREGABALIN

CAPSULE; ORAL

PREGABALIN

AB		<u>150MG</u>	<u>A205321 005</u>	Mar 29, 2023
AB		<u>200MG</u>	<u>A205321 006</u>	Mar 29, 2023
AB		<u>225MG</u>	<u>A205321 007</u>	Mar 29, 2023
AB		<u>300MG</u>	<u>A205321 008</u>	Mar 29, 2023
AB	CHANGZHOU PHARM	<u>25MG</u>	<u>A214322 008</u>	Sep 26, 2024
AB		<u>50MG</u>	<u>A214322 001</u>	Jul 15, 2021
AB		<u>75MG</u>	<u>A214322 002</u>	Jul 15, 2021
AB		<u>100MG</u>	<u>A214322 003</u>	Jul 15, 2021
AB		<u>150MG</u>	<u>A214322 004</u>	Jul 15, 2021
AB		<u>200MG</u>	<u>A214322 005</u>	Jul 15, 2021
AB		<u>225MG</u>	<u>A214322 006</u>	Jul 15, 2021
AB		<u>300MG</u>	<u>A214322 007</u>	Jul 15, 2021
AB	CHARTWELL RX	<u>50MG</u>	<u>A212865 001</u>	Mar 20, 2020
AB		<u>75MG</u>	<u>A212865 002</u>	Mar 20, 2020
AB		<u>100MG</u>	<u>A212865 003</u>	Mar 20, 2020
AB		<u>150MG</u>	<u>A212865 004</u>	Mar 20, 2020
AB	CREEKWOOD PHARMS	<u>25MG</u>	<u>A213423 001</u>	Mar 23, 2020
AB		<u>50MG</u>	<u>A213423 002</u>	Mar 23, 2020
AB		<u>75MG</u>	<u>A213423 003</u>	Mar 23, 2020
AB		<u>100MG</u>	<u>A213423 004</u>	Mar 23, 2020
AB		<u>150MG</u>	<u>A213423 005</u>	Mar 23, 2020
AB		<u>200MG</u>	<u>A213423 006</u>	Mar 23, 2020
AB		<u>225MG</u>	<u>A213423 007</u>	Mar 23, 2020
AB		<u>300MG</u>	<u>A213423 008</u>	Mar 23, 2020
AB	DR REDDYS	<u>25MG</u>	<u>A209664 001</u>	Jul 19, 2019
AB		<u>50MG</u>	<u>A209664 002</u>	Jul 19, 2019
AB		<u>75MG</u>	<u>A209664 003</u>	Jul 19, 2019
AB		<u>100MG</u>	<u>A209664 004</u>	Jul 19, 2019
AB		<u>150MG</u>	<u>A209664 005</u>	Jul 19, 2019
AB		<u>200MG</u>	<u>A209664 006</u>	Jul 19, 2019
AB		<u>225MG</u>	<u>A209664 007</u>	Jul 19, 2019
AB		<u>300MG</u>	<u>A209664 008</u>	Jul 19, 2019
AB	ESKAYEF	<u>25MG</u>	<u>A212988 001</u>	Mar 08, 2022
AB		<u>50MG</u>	<u>A212988 002</u>	Mar 08, 2022
AB		<u>75MG</u>	<u>A212988 003</u>	Mar 08, 2022
AB		<u>100MG</u>	<u>A212988 004</u>	Mar 08, 2022
AB		<u>150MG</u>	<u>A212988 005</u>	Mar 08, 2022
AB		<u>200MG</u>	<u>A212988 006</u>	Mar 08, 2022
AB		<u>225MG</u>	<u>A212988 007</u>	Mar 08, 2022
AB		<u>300MG</u>	<u>A212988 008</u>	Mar 08, 2022
AB	HETERO LABS LTD III	<u>25MG</u>	<u>A206912 001</u>	Oct 08, 2019
AB		<u>50MG</u>	<u>A206912 002</u>	Oct 08, 2019
AB		<u>75MG</u>	<u>A206912 003</u>	Oct 08, 2019
AB		<u>100MG</u>	<u>A206912 004</u>	Oct 08, 2019
AB		<u>150MG</u>	<u>A206912 005</u>	Oct 08, 2019
AB		<u>200MG</u>	<u>A206912 006</u>	Oct 08, 2019
AB		<u>225MG</u>	<u>A206912 007</u>	Oct 08, 2019
AB		<u>300MG</u>	<u>A206912 008</u>	Oct 08, 2019
AB	INVAGEN PHARMS	<u>25MG</u>	<u>A211384 001</u>	Jul 19, 2019
AB		<u>50MG</u>	<u>A211384 002</u>	Jul 19, 2019
AB		<u>75MG</u>	<u>A211384 003</u>	Jul 19, 2019
AB		<u>100MG</u>	<u>A211384 004</u>	Jul 19, 2019
AB		<u>150MG</u>	<u>A211384 005</u>	Jul 19, 2019
AB		<u>200MG</u>	<u>A211384 006</u>	Jul 19, 2019
AB		<u>225MG</u>	<u>A211384 007</u>	Jul 19, 2019
AB		<u>300MG</u>	<u>A211384 008</u>	Jul 19, 2019
AB	MACLEODS PHARMS LTD	<u>25MG</u>	<u>A205924 001</u>	Nov 12, 2024
AB		<u>50MG</u>	<u>A205924 002</u>	Nov 12, 2024
AB		<u>75MG</u>	<u>A205924 003</u>	Nov 12, 2024
AB		<u>100MG</u>	<u>A205924 004</u>	Nov 12, 2024
AB		<u>150MG</u>	<u>A205924 005</u>	Nov 12, 2024
AB		<u>200MG</u>	<u>A205924 006</u>	Nov 12, 2024
AB		<u>225MG</u>	<u>A205924 007</u>	Nov 12, 2024
AB		<u>300MG</u>	<u>A205924 008</u>	Nov 12, 2024
AB	MSN	<u>25MG</u>	<u>A209357 001</u>	Jul 19, 2019
AB		<u>50MG</u>	<u>A209357 002</u>	Jul 19, 2019
AB		<u>75MG</u>	<u>A209357 003</u>	Jul 19, 2019
AB		<u>100MG</u>	<u>A209357 004</u>	Jul 19, 2019
AB		<u>150MG</u>	<u>A209357 005</u>	Jul 19, 2019

PRESCRIPTION DRUG PRODUCT LIST

PREGABALIN

CAPSULE; ORAL

PREGABALIN

<u>AB</u>		<u>200MG</u>	<u>A209357 006</u>	Jul 19, 2019
<u>AB</u>		<u>225MG</u>	<u>A209357 007</u>	Jul 19, 2019
<u>AB</u>		<u>300MG</u>	<u>A209357 008</u>	Jul 19, 2019
<u>AB</u>	PRINSTON INC	<u>25MG</u>	<u>A207883 001</u>	Sep 01, 2022
<u>AB</u>		<u>50MG</u>	<u>A207883 002</u>	Sep 01, 2022
<u>AB</u>		<u>75MG</u>	<u>A207883 003</u>	Sep 01, 2022
<u>AB</u>		<u>100MG</u>	<u>A207883 004</u>	Sep 01, 2022
<u>AB</u>		<u>150MG</u>	<u>A207883 005</u>	Sep 01, 2022
<u>AB</u>		<u>200MG</u>	<u>A207883 006</u>	Sep 01, 2022
<u>AB</u>		<u>225MG</u>	<u>A207883 007</u>	Sep 01, 2022
<u>AB</u>		<u>300MG</u>	<u>A207883 008</u>	Sep 01, 2022
<u>AB</u>	RENATA	<u>50MG</u>	<u>A210585 001</u>	Dec 26, 2019
<u>AB</u>		<u>75MG</u>	<u>A210585 002</u>	Dec 26, 2019
<u>AB</u>		<u>100MG</u>	<u>A210585 003</u>	Dec 26, 2019
<u>AB</u>		<u>150MG</u>	<u>A210585 004</u>	Dec 26, 2019
<u>AB</u>		<u>200MG</u>	<u>A210585 005</u>	Dec 26, 2019
<u>AB</u>		<u>300MG</u>	<u>A210585 006</u>	Dec 26, 2019
<u>AB</u>	RISING	<u>25MG</u>	<u>A210432 001</u>	Jul 19, 2019
<u>AB</u>		<u>50MG</u>	<u>A210432 002</u>	Jul 19, 2019
<u>AB</u>		<u>75MG</u>	<u>A210432 003</u>	Jul 19, 2019
<u>AB</u>		<u>100MG</u>	<u>A210432 004</u>	Jul 19, 2019
<u>AB</u>		<u>150MG</u>	<u>A210432 005</u>	Jul 19, 2019
<u>AB</u>		<u>200MG</u>	<u>A210432 006</u>	Jul 19, 2019
<u>AB</u>		<u>225MG</u>	<u>A210432 007</u>	Jul 19, 2019
<u>AB</u>		<u>300MG</u>	<u>A210432 008</u>	Jul 19, 2019
<u>AB</u>	SCIEGEN PHARMS INC	<u>25MG</u>	<u>A208677 001</u>	Jul 19, 2019
<u>AB</u>		<u>50MG</u>	<u>A208677 002</u>	Jul 19, 2019
<u>AB</u>		<u>75MG</u>	<u>A208677 003</u>	Jul 19, 2019
<u>AB</u>		<u>100MG</u>	<u>A208677 004</u>	Jul 19, 2019
<u>AB</u>		<u>150MG</u>	<u>A208677 005</u>	Jul 19, 2019
<u>AB</u>		<u>200MG</u>	<u>A208677 006</u>	Jul 19, 2019
<u>AB</u>		<u>225MG</u>	<u>A208677 007</u>	Jul 19, 2019
<u>AB</u>		<u>300MG</u>	<u>A208677 008</u>	Jul 19, 2019
<u>AB</u>	STRIDES PHARMA	<u>25MG</u>	<u>A215755 001</u>	Apr 26, 2023
<u>AB</u>		<u>50MG</u>	<u>A215755 002</u>	Apr 26, 2023
<u>AB</u>		<u>75MG</u>	<u>A215755 003</u>	Apr 26, 2023
<u>AB</u>		<u>100MG</u>	<u>A215755 004</u>	Apr 26, 2023
<u>AB</u>		<u>150MG</u>	<u>A215755 005</u>	Apr 26, 2023
<u>AB</u>		<u>200MG</u>	<u>A215755 006</u>	Apr 26, 2023
<u>AB</u>		<u>225MG</u>	<u>A215755 007</u>	Apr 26, 2023
<u>AB</u>		<u>300MG</u>	<u>A215755 008</u>	Apr 26, 2023
<u>AB</u>	YILING	<u>75MG</u>	<u>A210891 001</u>	Sep 30, 2019
<u>AB</u>		<u>300MG</u>	<u>A210891 002</u>	Sep 30, 2019

SOLUTION; ORAL

LYRICA

<u>AA</u>	+!	UPJOHN	<u>20MG/ML</u>	<u>N022488 001</u>	Jan 04, 2010
-----------	----	--------	----------------	--------------------	--------------

PREGABALIN

<u>AA</u>		ALKEM LABS LTD	<u>20MG/ML</u>	<u>A207623 001</u>	Jul 19, 2019
<u>AA</u>		ANDA REPOSITORY	<u>20MG/ML</u>	<u>A212604 001</u>	Feb 18, 2022

TABLET, EXTENDED RELEASE; ORAL

LYRICA CR

<u>AB</u>	+	UPJOHN	<u>82.5MG</u>	<u>N209501 001</u>	Oct 11, 2017
<u>AB</u>	+		<u>165MG</u>	<u>N209501 002</u>	Oct 11, 2017
<u>AB</u>	+!		<u>330MG</u>	<u>N209501 003</u>	Oct 11, 2017

PREGABALIN

<u>AB</u>		ALVOGEN	<u>82.5MG</u>	<u>A211593 001</u>	Apr 13, 2021
<u>AB</u>			<u>165MG</u>	<u>A211593 002</u>	Apr 13, 2021
<u>AB</u>			<u>330MG</u>	<u>A211593 003</u>	Apr 13, 2021
<u>AB</u>		APOTEX	<u>165MG</u>	<u>A213313 001</u>	Apr 13, 2021
<u>AB</u>			<u>330MG</u>	<u>A213313 002</u>	Apr 13, 2021
<u>AB</u>		EPIC PHARMA LLC	<u>82.5MG</u>	<u>A214496 001</u>	Jun 01, 2023
<u>AB</u>			<u>165MG</u>	<u>A214496 002</u>	Jun 01, 2023
<u>AB</u>			<u>330MG</u>	<u>A214496 003</u>	Jun 01, 2023
<u>AB</u>		MSN	<u>82.5MG</u>	<u>A213226 001</u>	Apr 13, 2021
<u>AB</u>			<u>165MG</u>	<u>A213226 002</u>	Apr 13, 2021
<u>AB</u>			<u>330MG</u>	<u>A213226 003</u>	Apr 13, 2021
<u>AB</u>		RUBICON	<u>82.5MG</u>	<u>A215249 001</u>	Mar 22, 2022
<u>AB</u>			<u>165MG</u>	<u>A215249 002</u>	Mar 22, 2022
<u>AB</u>			<u>330MG</u>	<u>A215249 003</u>	Mar 22, 2022

PRESCRIPTION DRUG PRODUCT LIST

PRETOMANID

TABLET; ORAL

PRETOMANID

+! MYLAN IRELAND LTD 200MG N212862 001 Aug 14, 2019

PRILOCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

PRILOCAINE HYDROCHLORIDE

! SEPTODONT INC 4% A079235 001 Sep 29, 2010

PRIMAQUINE PHOSPHATE

TABLET; ORAL

PRIMAQUINE**AB** +! SANOFI AVENTIS US **EQ 15MG BASE** **N008316 001****PRIMAQUINE PHOSPHATE****AB** NOVAST LABS **EQ 15MG BASE** **A206043 001** Jun 23, 2016**AB** UNICHEM **EQ 15MG BASE** **A204476 001** Feb 25, 2014PRIMIDONE

TABLET; ORAL

MYSOLINE**AB** +! VALEANT **50MG** **N009170 003****AB** + **250MG** **N009170 002****PRIMIDONE****AB** AMNEAL PHARM **50MG** **A040866 001** Apr 23, 2008**AB** **250MG** **A040866 002** Apr 23, 2008**AB** ANDA REPOSITORY **50MG** **A040626 001** Sep 29, 2005**AB** **250MG** **A040626 002** Sep 29, 2005**AB** CARNEGIE **50MG** **A218366 001** Jan 23, 2024**AB** **250MG** **A218366 002** Jan 23, 2024**AB** LANNETT **50MG** **A084903 002** May 24, 2001**AB** **250MG** **A084903 001****AB** OXFORD PHARMS **50MG** **A040586 001** Feb 24, 2005**AB** **250MG** **A040586 002** Feb 24, 2005**AB** RUBICON **50MG** **A214896 001** Jun 28, 2022**AB** **250MG** **A214896 002** Jun 28, 2022**AB** WATSON LABS **250MG** **A083551 001**

RUBICON 125MG A214896 003 Dec 28, 2022

PROBENECID

TABLET; ORAL

PROBALAN**AB** LANNETT **500MG** **A080966 001****PROBENECID****AB** RISING **500MG** **A217020 001** Nov 20, 2023**AB** ! WATSON LABS TEVA **500MG** **A084442 004** Mar 29, 1983PROBENECID; SULOPENEM ETZADROXIL

TABLET; ORAL

ORLYNVAH

+! ITERUM THERAP 500MG; 500MG N213972 001 Oct 25, 2024

PROCAINAMIDE HYDROCHLORIDE

INJECTABLE; INJECTION

PROCAINAMIDE HYDROCHLORIDE**AP** GLAND PHARMA LTD **100MG/ML** **A218135 001** Jul 22, 2024**AP** **500MG/ML** **A218135 002** Jul 22, 2024**AP** ! HOSPIRA **100MG/ML** **A089069 001** Feb 12, 1986**AP** ! **500MG/ML** **A089070 001** Feb 12, 1986**AP** INTL MEDICATION **100MG/ML** **A088636 001** Jul 31, 1984**AP** NEXUS **100MG/ML** **A206332 001** Oct 13, 2017**AP** **500MG/ML** **A206332 002** Oct 13, 2017PROCARBAZINE HYDROCHLORIDE

CAPSULE; ORAL

MATULANE

+! LEADIANT BIOSCI INC EQ 50MG BASE N016785 001

PROCHLORPERAZINE

SUPPOSITORY; RECTAL

COMPRO**AB** PADAGIS US **25MG** **A040246 001** Jun 28, 2000**PROCHLORPERAZINE****AB** ! COSETTE **25MG** **A040058 001** Nov 24, 1993

PRESCRIPTION DRUG PRODUCT LIST

PROCHLORPERAZINE EDISYLATE

INJECTABLE; INJECTION

PROCHLORPERAZINE EDISYLATE

<u>AP</u>	AMNEAL	<u>EQ 5MG BASE/ML</u>	<u>A214192 001</u>	Nov 28, 2022
<u>AP</u>	! AVET LIFESCIENCES	<u>EQ 5MG BASE/ML</u>	<u>A204147 001</u>	Oct 15, 2013
<u>AP</u>	CAPLIN	<u>EQ 5MG BASE/ML</u>	<u>A214379 001</u>	Apr 22, 2021
<u>AP</u>	EUGIA PHARMA	<u>EQ 5MG BASE/ML</u>	<u>A213873 001</u>	Jul 14, 2022
<u>AP</u>	GLAND PHARMA LTD	<u>EQ 5MG BASE/ML</u>	<u>A214107 001</u>	Sep 22, 2021
<u>AP</u>	HIKMA	<u>EQ 5MG BASE/ML</u>	<u>A089903 001</u>	Aug 29, 1989
<u>AP</u>	MYLAN LABS LTD	<u>EQ 5MG BASE/ML</u>	<u>A210710 001</u>	Oct 25, 2018
<u>AP</u>	NEXUS	<u>EQ 5MG BASE/ML</u>	<u>A204860 001</u>	Feb 15, 2019
<u>AP</u>	SAGENT	<u>EQ 5MG BASE/ML</u>	<u>A040540 001</u>	May 28, 2004
<u>AP</u>	SOMERSET THERAPS LLC	<u>EQ 5MG BASE/ML</u>	<u>A212257 001</u>	Sep 13, 2024
<u>AP</u>	VIWIT PHARM	<u>EQ 5MG BASE/ML</u>	<u>A213626 001</u>	Sep 28, 2021

PROCHLORPERAZINE MALEATE

TABLET; ORAL

PROCHLORPERAZINE MALEATE

<u>AB</u>	AJANTA PHARMA LTD	<u>EQ 5MG BASE</u>	<u>A218515 001</u>	May 09, 2024
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A218515 002</u>	May 09, 2024
<u>AB</u>	AMNEAL	<u>EQ 5MG BASE</u>	<u>A216598 001</u>	Apr 17, 2023
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A216598 002</u>	Apr 17, 2023
<u>AB</u>	BIONPHARMA	<u>EQ 5MG BASE</u>	<u>A217478 001</u>	Apr 04, 2023
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A217478 002</u>	Apr 04, 2023
<u>AB</u>	GLENMARK PHARMS LTD	<u>EQ 5MG BASE</u>	<u>A216595 001</u>	Mar 17, 2023
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A216595 002</u>	Mar 17, 2023
<u>AB</u>	LEADING	<u>EQ 5MG BASE</u>	<u>A218912 001</u>	Sep 30, 2024
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A218912 002</u>	Sep 30, 2024
<u>AB</u>	NOVITIUM PHARMA	<u>EQ 5MG BASE</u>	<u>A216202 001</u>	Jun 13, 2022
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A216202 002</u>	Jun 13, 2022
<u>AB</u>	ZYDUS LIFESCIENCES	<u>EQ 5MG BASE</u>	<u>A216495 001</u>	Aug 08, 2022
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A216495 002</u>	Aug 08, 2022
<u>PROCOMP</u>				
<u>AB</u>	JUBILANT CADISTA	<u>EQ 5MG BASE</u>	<u>A040268 001</u>	Feb 27, 1998
<u>AB</u>	!	<u>EQ 10MG BASE</u>	<u>A040268 002</u>	Feb 27, 1998

PROGESTERONE

CAPSULE; ORAL

PROGESTERONE

<u>AB</u>	AMNEAL PHARMS NY	<u>100MG</u>	<u>A207724 001</u>	Sep 07, 2017
<u>AB</u>		<u>200MG</u>	<u>A207724 002</u>	Sep 07, 2017
<u>AB</u>	BIONPHARMA	<u>100MG</u>	<u>A200900 001</u>	Aug 16, 2013
<u>AB</u>		<u>200MG</u>	<u>A200900 002</u>	Aug 16, 2013
<u>AB</u>	DR REDDYS	<u>100MG</u>	<u>A208801 001</u>	Feb 28, 2017
<u>AB</u>		<u>200MG</u>	<u>A208801 002</u>	Feb 28, 2017
<u>AB</u>	EUGIA PHARMA	<u>100MG</u>	<u>A211285 001</u>	Oct 26, 2018
<u>AB</u>		<u>200MG</u>	<u>A211285 002</u>	Oct 26, 2018
<u>AB</u>	SOFGEN PHARMS	<u>100MG</u>	<u>A200456 001</u>	Sep 28, 2012
<u>AB</u>		<u>200MG</u>	<u>A200456 002</u>	Sep 28, 2012
<u>AB</u>	XIROMED	<u>100MG</u>	<u>A205229 001</u>	Oct 20, 2017
<u>AB</u>		<u>200MG</u>	<u>A205229 002</u>	Oct 20, 2017

PROMETRIUM

<u>AB</u>	+ VIRTUS	<u>100MG</u>	<u>N019781 001</u>	May 14, 1998
<u>AB</u>	+	<u>200MG</u>	<u>N019781 002</u>	Oct 15, 1999

GEL; VAGINAL

CRINONE

+	!	ABBVIE	4%	N020701 001	Jul 31, 1997
+	!		8%	N020701 002	Jul 31, 1997

INJECTABLE; INJECTION

PROGESTERONE

<u>AO</u>	!	EUGIA PHARMA	<u>50MG/ML</u>	<u>A210965 001</u>	Dec 06, 2018
<u>AO</u>		FRESENIUS KABI USA	<u>50MG/ML</u>	<u>A075906 001</u>	Apr 25, 2001
<u>AO</u>		HIKMA FARMACEUTICA	<u>50MG/ML</u>	<u>A091033 001</u>	Oct 28, 2010
<u>AO</u>		XIROMED	<u>50MG/ML</u>	<u>A215634 001</u>	Jan 05, 2022

INSERT; VAGINAL

ENDOMETRIN

+	!	FERRING	100MG	N022057 001	Jun 21, 2007
---	---	---------	-------	-------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

PROMETHAZINE HYDROCHLORIDE

INJECTABLE; INJECTION

PROMETHAZINE HYDROCHLORIDE

<u>AP</u>	<u>+</u> !	HIKMA	<u>25MG/ML</u>	<u>A083312</u>	<u>001</u>	
<u>AP</u>	<u>+</u> !		<u>50MG/ML</u>	<u>A083312</u>	<u>002</u>	
<u>AP</u>		XGEN PHARMS	<u>25MG/ML</u>	<u>A040737</u>	<u>001</u>	Apr 24, 2008
<u>AP</u>			<u>50MG/ML</u>	<u>A040737</u>	<u>002</u>	Apr 24, 2008

SUPPOSITORY; RECTAL

PROMETHAZINE HYDROCHLORIDE

<u>AB</u>		ANNORA PHARMA	<u>12.5MG</u>	<u>A216446</u>	<u>001</u>	Nov 02, 2022
<u>AB</u>			<u>25MG</u>	<u>A216446</u>	<u>002</u>	Nov 02, 2022
<u>AB</u>		COSETTE	<u>12.5MG</u>	<u>A040428</u>	<u>002</u>	Mar 31, 2003
<u>AB</u>	<u>!</u>		<u>25MG</u>	<u>A040428</u>	<u>001</u>	Feb 05, 2002
<u>AB</u>		PADAGIS ISRAEL	<u>12.5MG</u>	<u>A040500</u>	<u>001</u>	Jun 30, 2003
<u>AB</u>			<u>25MG</u>	<u>A040500</u>	<u>002</u>	Jun 30, 2003
<u>AB</u>		TARO	<u>12.5MG</u>	<u>A040603</u>	<u>001</u>	Oct 26, 2006
<u>AB</u>			<u>25MG</u>	<u>A040603</u>	<u>002</u>	Oct 26, 2006
		PROMETHEGAN				
	<u>!</u>	COSETTE	50MG	A087165	001	Aug 14, 1987

SYRUP; ORAL

PROMETHAZINE HYDROCHLORIDE

<u>AA</u>		CHARTWELL RX	<u>6.25MG/5ML</u>	<u>A040026</u>	<u>001</u>	Sep 25, 1998
<u>AA</u>		NOSTRUM LABS INC	<u>6.25MG/5ML</u>	<u>A040891</u>	<u>001</u>	Mar 13, 2009
<u>AA</u>		QUAGEN	<u>6.25MG/5ML</u>	<u>A213890</u>	<u>001</u>	Jul 12, 2021
<u>AA</u>		TARO	<u>6.25MG/5ML</u>	<u>A040718</u>	<u>001</u>	Apr 04, 2007
<u>AA</u>		TRIS PHARMA INC	<u>6.25MG/5ML</u>	<u>A091675</u>	<u>001</u>	Jun 28, 2012

PROMETHAZINE PLAIN

<u>AA</u>	<u>+</u> !	WOCKHARDT BIO AG	<u>6.25MG/5ML</u>	<u>A087953</u>	<u>001</u>	Nov 15, 1982
-----------	------------	------------------	-------------------	----------------	------------	--------------

TABLET; ORAL

PROMETHAZINE HYDROCHLORIDE

<u>AB</u>		AMNEAL PHARMS NY	<u>12.5MG</u>	<u>A091179</u>	<u>001</u>	Dec 13, 2010
<u>AB</u>			<u>25MG</u>	<u>A091179</u>	<u>002</u>	Dec 13, 2010
<u>AB</u>			<u>50MG</u>	<u>A091179</u>	<u>003</u>	Dec 13, 2010
<u>AB</u>		KVK TECH	<u>12.5MG</u>	<u>A040712</u>	<u>002</u>	May 04, 2007
<u>AB</u>			<u>25MG</u>	<u>A040712</u>	<u>001</u>	Jul 31, 2006
<u>AB</u>			<u>50MG</u>	<u>A040712</u>	<u>003</u>	Jul 31, 2006
<u>AB</u>		PRINSTON INC	<u>12.5MG</u>	<u>A040622</u>	<u>001</u>	Jul 18, 2006
<u>AB</u>			<u>25MG</u>	<u>A040622</u>	<u>002</u>	Jul 18, 2006
<u>AB</u>			<u>50MG</u>	<u>A040622</u>	<u>003</u>	Jul 18, 2006
<u>AB</u>		QUAGEN	<u>12.5MG</u>	<u>A040673</u>	<u>001</u>	Mar 05, 2008
<u>AB</u>			<u>25MG</u>	<u>A040673</u>	<u>002</u>	Mar 05, 2008
<u>AB</u>			<u>50MG</u>	<u>A040673</u>	<u>003</u>	Mar 05, 2008
<u>AB</u>	<u>+</u>	SANDOZ	<u>25MG</u>	<u>A084176</u>	<u>003</u>	
<u>AB</u>	<u>+</u> !		<u>50MG</u>	<u>A084176</u>	<u>001</u>	
<u>AB</u>		STRIDES PHARMA	<u>12.5MG</u>	<u>A209177</u>	<u>001</u>	Jun 30, 2017
<u>AB</u>			<u>25MG</u>	<u>A209177</u>	<u>002</u>	Jun 30, 2017
<u>AB</u>			<u>50MG</u>	<u>A209177</u>	<u>003</u>	Jun 30, 2017
<u>AB</u>		WATSON LABS	<u>25MG</u>	<u>A083426</u>	<u>001</u>	
<u>AB</u>			<u>50MG</u>	<u>A083711</u>	<u>001</u>	
<u>AB</u>		ZYDUS PHARMS USA	<u>12.5MG</u>	<u>A040596</u>	<u>001</u>	Nov 18, 2005
<u>AB</u>			<u>25MG</u>	<u>A040596</u>	<u>002</u>	Nov 18, 2005
<u>AB</u>			<u>50MG</u>	<u>A040596</u>	<u>003</u>	Nov 18, 2005

PROPAFENONE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

PROPAFENONE HYDROCHLORIDE

<u>AB</u>		AUROBINDO PHARMA LTD	<u>225MG</u>	<u>A213096</u>	<u>001</u>	Feb 21, 2023
<u>AB</u>			<u>325MG</u>	<u>A213096</u>	<u>002</u>	Feb 21, 2023
<u>AB</u>			<u>425MG</u>	<u>A213096</u>	<u>003</u>	Feb 21, 2023
<u>AB</u>		GLENMARK PHARMS LTD	<u>225MG</u>	<u>A205268</u>	<u>001</u>	Sep 08, 2017
<u>AB</u>			<u>325MG</u>	<u>A205268</u>	<u>002</u>	Sep 08, 2017
<u>AB</u>			<u>425MG</u>	<u>A205268</u>	<u>003</u>	Sep 08, 2017
<u>AB</u>		RISING	<u>225MG</u>	<u>A205956</u>	<u>001</u>	Jul 02, 2018
<u>AB</u>			<u>325MG</u>	<u>A205956</u>	<u>002</u>	Jul 02, 2018
<u>AB</u>			<u>425MG</u>	<u>A205956</u>	<u>003</u>	Jul 02, 2018
<u>AB</u>		SINOTHERAPEUTICS INC	<u>225MG</u>	<u>A210339</u>	<u>001</u>	Jan 04, 2019
<u>AB</u>			<u>325MG</u>	<u>A210339</u>	<u>002</u>	Jan 04, 2019
<u>AB</u>			<u>425MG</u>	<u>A210339</u>	<u>003</u>	Jan 04, 2019
<u>AB</u>		STRIDES PHARMA	<u>225MG</u>	<u>A078540</u>	<u>001</u>	Oct 18, 2010
<u>AB</u>			<u>325MG</u>	<u>A078540</u>	<u>002</u>	Oct 18, 2010
<u>AB</u>	<u>!</u>		<u>425MG</u>	<u>A078540</u>	<u>003</u>	Oct 18, 2010

PRESCRIPTION DRUG PRODUCT LIST

PROPAFENONE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE;ORAL

PROPAFENONE HYDROCHLORIDE

<u>AB</u>	UPSHER SMITH LABS	<u>225MG</u>	<u>A212744 001</u>	Jun 25, 2020
<u>AB</u>		<u>325MG</u>	<u>A212744 002</u>	Jun 25, 2020
<u>AB</u>		<u>425MG</u>	<u>A212744 003</u>	Jun 25, 2020
<u>AB</u>	WATSON LABS INC	<u>225MG</u>	<u>A202688 001</u>	Aug 24, 2015
<u>AB</u>		<u>325MG</u>	<u>A202688 002</u>	Aug 24, 2015
<u>AB</u>		<u>425MG</u>	<u>A202688 003</u>	Aug 24, 2015
<u>AB</u>	ZYDUS LIFESCIENCES	<u>225MG</u>	<u>A214184 001</u>	Apr 21, 2021
<u>AB</u>		<u>325MG</u>	<u>A214184 002</u>	Apr 21, 2021
<u>AB</u>		<u>425MG</u>	<u>A214184 003</u>	Apr 21, 2021

TABLET;ORAL

PROPAFENONE HYDROCHLORIDE

<u>AB</u>	ANI PHARMS	<u>150MG</u>	<u>A076550 001</u>	Apr 23, 2004
<u>AB</u>		<u>225MG</u>	<u>A076550 002</u>	Apr 23, 2004
<u>AB</u>		<u>300MG</u>	<u>A076550 003</u>	Apr 23, 2004
<u>AB</u>	AUROBINDO PHARMA	<u>150MG</u>	<u>A202445 001</u>	May 11, 2016
<u>AB</u>		<u>225MG</u>	<u>A202445 002</u>	May 11, 2016
<u>AB</u>		<u>300MG</u>	<u>A202445 003</u>	May 11, 2016
<u>AB</u>	STRIDES PHARMA	<u>150MG</u>	<u>A075938 001</u>	Oct 17, 2002
<u>AB</u>		<u>225MG</u>	<u>A075938 002</u>	Oct 17, 2002
<u>AB</u>	!	<u>300MG</u>	<u>A075938 003</u>	Oct 17, 2002
<u>AB</u>	WATSON LABS	<u>150MG</u>	<u>A075203 001</u>	Oct 24, 2000
<u>AB</u>		<u>225MG</u>	<u>A075203 002</u>	Oct 24, 2000

PROPARACAINE HYDROCHLORIDE

SOLUTION/DROPS;OPHTHALMIC

ALCAINE

<u>AT</u>	ALCON LABS INC	<u>0.5%</u>	<u>A080027 001</u>	
-----------	----------------	-------------	--------------------	--

PROPARACAINE HYDROCHLORIDE

<u>AT</u>	!	BAUSCH AND LOMB	<u>0.5%</u>	<u>A040074 001</u>	Sep 29, 1995
<u>AT</u>		RISING	<u>0.5%</u>	<u>A040277 001</u>	Mar 16, 2000
<u>AT</u>		SOMERSET THERAPS LLC	<u>0.5%</u>	<u>A215816 001</u>	Aug 09, 2024

PROPOFOL

INJECTABLE;INJECTION

DIPRIVAN

<u>AB</u>	+	FRESENIUS KABI USA	<u>10MG/ML</u>	<u>N019627 002</u>	Jun 11, 1996
-----------	---	--------------------	----------------	--------------------	--------------

PROPOFOL

<u>AB</u>		AMNEAL	<u>10MG/ML</u>	<u>A217525 001</u>	Aug 15, 2024
<u>AB</u>		ASPIRO	<u>10MG/ML</u>	<u>A217013 001</u>	Jun 20, 2024
<u>AB</u>		AVET LIFESCIENCES	<u>10MG/ML</u>	<u>A206408 001</u>	Oct 12, 2021
<u>AB</u>		DR REDDYS	<u>10MG/ML</u>	<u>A205067 001</u>	Nov 15, 2018
<u>AB</u>		HIKMA	<u>10MG/ML</u>	<u>A074848 001</u>	Apr 19, 2005
<u>AB</u>		HOSPIRA	<u>10MG/ML</u>	<u>A077908 001</u>	Mar 17, 2006
<u>AB</u>		INNOPHARMA	<u>10MG/ML</u>	<u>A205576 001</u>	Sep 16, 2020
<u>AB</u>		SAGENT PHARMS INC	<u>10MG/ML</u>	<u>A075102 001</u>	Jan 04, 1999

PROPRANOLOL HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE;ORAL

INDERAL LA

<u>AB</u>	+	ANI PHARMS	<u>60MG</u>	<u>N018553 004</u>	Mar 18, 1987
<u>AB</u>	+		<u>80MG</u>	<u>N018553 002</u>	Apr 19, 1983
<u>AB</u>	+		<u>120MG</u>	<u>N018553 003</u>	Apr 19, 1983
<u>AB</u>	+		<u>160MG</u>	<u>N018553 001</u>	Apr 19, 1983

PROPRANOLOL HYDROCHLORIDE

<u>AB</u>		ACTAVIS ELIZABETH	<u>60MG</u>	<u>A078494 001</u>	Aug 10, 2007
<u>AB</u>			<u>80MG</u>	<u>A078494 002</u>	Aug 10, 2007
<u>AB</u>			<u>120MG</u>	<u>A078494 003</u>	Aug 10, 2007
<u>AB</u>			<u>160MG</u>	<u>A078494 004</u>	Aug 10, 2007
<u>AB</u>		ADARE PHARMS INC	<u>60MG</u>	<u>A078703 001</u>	Jul 15, 2011
<u>AB</u>			<u>80MG</u>	<u>A078703 002</u>	Jul 15, 2011
<u>AB</u>			<u>120MG</u>	<u>A078703 003</u>	Jul 15, 2011
<u>AB</u>			<u>160MG</u>	<u>A078703 004</u>	Jul 15, 2011
<u>AB</u>		AMTA	<u>60MG</u>	<u>A212026 001</u>	Jan 06, 2020
<u>AB</u>			<u>80MG</u>	<u>A212026 002</u>	Jan 06, 2020
<u>AB</u>			<u>120MG</u>	<u>A212026 003</u>	Jan 06, 2020
<u>AB</u>			<u>160MG</u>	<u>A212026 004</u>	Jan 06, 2020
<u>AB</u>		NORTEC DEV ASSOC	<u>60MG</u>	<u>A078065 001</u>	Jan 26, 2007
<u>AB</u>			<u>80MG</u>	<u>A078065 002</u>	Jan 26, 2007
<u>AB</u>			<u>120MG</u>	<u>A078065 003</u>	Jan 26, 2007
<u>AB</u>			<u>160MG</u>	<u>A078065 004</u>	Jan 26, 2007

PRESCRIPTION DRUG PRODUCT LIST

PROPRANOLOL HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

PROPRANOLOL HYDROCHLORIDE

AB	ZYDUS PHARMS USA INC	60MG	A090321 001	Mar 25, 2011
AB		80MG	A090321 002	Mar 25, 2011
AB		120MG	A090321 003	Mar 25, 2011
AB		160MG	A090321 004	Mar 25, 2011
	INNOPRAN XL			
EX +	ANI PHARMS	80MG	N021438 001	Mar 12, 2003
EX +!		120MG	N021438 002	Mar 12, 2003

INJECTABLE; INJECTION

PROPRANOLOL HYDROCHLORIDE

AP	FRESENIUS KABI USA	1MG/ML	A075826 001	Aug 31, 2001
AP !	HIKMA FARMACEUTICA	1MG/ML	A077760 001	Jan 31, 2008
	SOLUTION; ORAL			
	HEMANGEOL			
	+! PIERRE FABRE	4.28MG/ML	N205410 001	Mar 14, 2014
	PROPRANOLOL HYDROCHLORIDE			
	! HIKMA	20MG/5ML	A070979 001	May 15, 1987
	! HIKMA	40MG/5ML	A070690 001	May 15, 1987

TABLET; ORAL

PROPRANOLOL HYDROCHLORIDE

AB	ENDO OPERATIONS	10MG	A070221 002	Aug 01, 1986
AB		20MG	A070221 003	Aug 01, 1986
AB		40MG	A070221 004	Aug 01, 1986
AB		60MG	A070221 005	Sep 24, 1986
AB		80MG	A070221 001	Apr 14, 1986
AB	IMPAX LABS INC	10MG	A071972 001	Apr 06, 1988
AB		20MG	A071972 002	Apr 06, 1988
AB		40MG	A071972 003	Apr 06, 1988
AB		60MG	A071976 002	May 13, 1986
AB !		80MG	A071976 001	Apr 06, 1988
AB	INNOGENIX	10MG	A070322 002	Oct 22, 1985
AB		20MG	A070322 003	Oct 22, 1985
AB		40MG	A070322 004	Oct 22, 1985
AB		60MG	A070322 005	Sep 24, 1986
AB		80MG	A070322 001	Aug 04, 1986
AB	IPCA LABS LTD	10MG	A078955 001	Jun 02, 2008
AB		20MG	A078955 002	Jun 02, 2008
AB		40MG	A078955 003	Jun 02, 2008
AB		60MG	A078955 004	Jun 02, 2008
AB		80MG	A078955 005	Jun 02, 2008
AB	MYLAN	10MG	A070213 002	Nov 19, 1985
AB		20MG	A070213 003	Nov 19, 1985
AB		40MG	A070213 001	Nov 19, 1985
AB		60MG	A070213 005	Apr 08, 2011
AB		80MG	A070213 004	Nov 19, 1985
AB	NORTHSTAR HLTHCARE	10MG	A078213 001	Jan 10, 2008
AB		20MG	A078213 002	Jan 10, 2008
AB		40MG	A078213 003	Jan 10, 2008
AB		60MG	A078213 004	Jan 10, 2008
AB		80MG	A078213 005	Jan 10, 2008
AB	WATSON LABS	10MG	A070175 001	May 13, 1986
AB		20MG	A070176 001	May 13, 1986
AB		40MG	A070177 001	May 13, 1986
AB		80MG	A070178 001	May 13, 1986

PROPYLTHIOURACIL

TABLET; ORAL

PROPYLTHIOURACIL

AB +!	ENDO OPERATIONS	50MG	N006188 001	
AB	MACLEODS PHARMS LTD	50MG	A208867 001	May 10, 2023
BD	ACTAVIS ELIZABETH	50MG	A080172 001	
BD	QUAGEN	50MG	A080154 001	

PROTAMINE SULFATE

INJECTABLE; INJECTION

PROTAMINE SULFATE

	! FRESENIUS KABI USA	10MG/ML	A089454 001	Apr 07, 1987
--	----------------------	---------	-------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

PROTRIPTYLINE HYDROCHLORIDE

TABLET; ORAL

PROTRIPTYLINE HYDROCHLORIDE

<u>AB</u>	EPIC PHARMA LLC	<u>5MG</u>	<u>A202220</u>	<u>001</u>	Nov 19, 2012
<u>AB</u>		<u>10MG</u>	<u>A202220</u>	<u>002</u>	Nov 19, 2012
<u>AB</u>	HIKMA	<u>5MG</u>	<u>A078913</u>	<u>001</u>	Sep 16, 2008
<u>AB</u>	!	<u>10MG</u>	<u>A078913</u>	<u>002</u>	Sep 16, 2008
<u>AB</u>	SIGMAPHARM LABS LLC	<u>5MG</u>	<u>A090462</u>	<u>001</u>	May 03, 2010
<u>AB</u>		<u>10MG</u>	<u>A090462</u>	<u>002</u>	May 03, 2010

PRUCALOPRIDE SUCCINATE

TABLET; ORAL

MOTEGRITY

<u>AB</u>	+	TAKEDA PHARMS USA	<u>EQ 1MG BASE</u>	<u>N210166</u>	<u>001</u>	Dec 14, 2018
<u>AB</u>	+	!	<u>EQ 2MG BASE</u>	<u>N210166</u>	<u>002</u>	Dec 14, 2018

PRUCALOPRIDE SUCCINATE

<u>AB</u>	NOVITIUM PHARMA	<u>EQ 1MG BASE</u>	<u>A218492</u>	<u>001</u>	Dec 26, 2024
<u>AB</u>		<u>EQ 2MG BASE</u>	<u>A218492</u>	<u>002</u>	Dec 26, 2024

PYRAZINAMIDE

TABLET; ORAL

PYRAZINAMIDE

<u>AB</u>	!	HIKMA	<u>500MG</u>	<u>A081319</u>	<u>001</u>	Jun 30, 1992
<u>AB</u>		MACLEODS PHARMS LTD	<u>500MG</u>	<u>A212541</u>	<u>001</u>	Jul 27, 2020
<u>AB</u>	+	NOVITIUM PHARMA	<u>500MG</u>	<u>A080157</u>	<u>001</u>	

PYRIDOSTIGMINE BROMIDE

INJECTABLE; INJECTION

MESTINON

<u>AP</u>	+	!	BAUSCH	<u>5MG/ML</u>	<u>N009830</u>	<u>001</u>
-----------	---	---	--------	---------------	----------------	------------

REGONOL

<u>AP</u>		SANDOZ	<u>5MG/ML</u>	<u>N017398</u>	<u>001</u>	
-----------	--	--------	---------------	----------------	------------	--

SYRUP; ORAL

MESTINON

<u>AA</u>	+	!	BAUSCH	<u>60MG/5ML</u>	<u>N015193</u>	<u>001</u>
-----------	---	---	--------	-----------------	----------------	------------

PYRIDOSTIGMINE BROMIDE

<u>AA</u>		AMNEAL	<u>60MG/5ML</u>	<u>A212702</u>	<u>001</u>	Jan 10, 2020
<u>AA</u>		MILLA PHARMS	<u>60MG/5ML</u>	<u>A212405</u>	<u>001</u>	Apr 19, 2022
<u>AA</u>		MSN	<u>60MG/5ML</u>	<u>A216512</u>	<u>001</u>	Sep 21, 2023
<u>AA</u>		NOVITIUM PHARMA	<u>60MG/5ML</u>	<u>A211694</u>	<u>001</u>	Mar 08, 2019
<u>AA</u>		RISING	<u>60MG/5ML</u>	<u>A208797</u>	<u>001</u>	Jan 09, 2020

TABLET; ORAL

MESTINON

<u>AB</u>	+	!	BAUSCH	<u>60MG</u>	<u>N009829</u>	<u>002</u>
-----------	---	---	--------	-------------	----------------	------------

PYRIDOSTIGMINE BROMIDE

<u>AB</u>		ANI PHARMS	<u>60MG</u>	<u>A040512</u>	<u>001</u>	Oct 08, 2003
<u>AB</u>		IMPAX LABS	<u>30MG</u>	<u>A040502</u>	<u>002</u>	Jun 28, 2022
<u>AB</u>			<u>60MG</u>	<u>A040502</u>	<u>001</u>	Apr 24, 2003
<u>AB</u>		MLV	<u>30MG</u>	<u>A211181</u>	<u>002</u>	May 14, 2019
<u>AB</u>			<u>60MG</u>	<u>A211181</u>	<u>001</u>	Jul 20, 2018
<u>AB</u>		ZYDUS PHARMS	<u>60MG</u>	<u>A205650</u>	<u>001</u>	Jun 22, 2015

TABLET, EXTENDED RELEASE; ORAL

MESTINON

<u>AB</u>	+	!	BAUSCH	<u>180MG</u>	<u>N011665</u>	<u>001</u>
-----------	---	---	--------	--------------	----------------	------------

PYRIDOSTIGMINE BROMIDE

<u>AB</u>		ALVOGEN	<u>180MG</u>	<u>A204737</u>	<u>001</u>	Jun 26, 2015
<u>AB</u>		IMPAX LABS INC	<u>180MG</u>	<u>A203184</u>	<u>001</u>	Sep 15, 2015
<u>AB</u>		RISING	<u>180MG</u>	<u>A205464</u>	<u>001</u>	Aug 15, 2017

PYRIDOXINE HYDROCHLORIDE

INJECTABLE; INJECTION

PYRIDOXINE HYDROCHLORIDE

+	!	FRESENIUS KABI USA	100MG/ML	A080618	001
---	---	--------------------	----------	---------	-----

PYRIMETHAMINE

TABLET; ORAL

DARAPRIM

<u>AB</u>	+	!	TILDE SCIENCES	<u>25MG</u>	<u>N008578</u>	<u>001</u>
-----------	---	---	----------------	-------------	----------------	------------

PYRIMETHAMINE

<u>AB</u>		ALVOGEN	<u>25MG</u>	<u>A211271</u>	<u>001</u>	Jul 27, 2021
<u>AB</u>		AUROBINDO PHARMA LTD	<u>25MG</u>	<u>A216983</u>	<u>001</u>	Oct 25, 2022
<u>AB</u>		SANALUZ	<u>25MG</u>	<u>A207127</u>	<u>001</u>	Feb 28, 2020
<u>AB</u>		TEVA PHARMS	<u>25MG</u>	<u>A215506</u>	<u>001</u>	Aug 13, 2021

PRESCRIPTION DRUG PRODUCT LIST

QUAZEPAM

TABLET; ORAL

DORAL

+! GALT PHARMS

15MG

N018708 001 Dec 27, 1985

QUETIAPINE FUMARATE

TABLET; ORAL

QUETIAPINE FUMARATE

<u>AB</u>	ACCORD HLTHCARE	<u>EQ 25MG BASE</u>	<u>A202152 001</u>	Mar 27, 2012
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A202152 002</u>	Mar 27, 2012
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A202152 003</u>	Mar 27, 2012
<u>AB</u>		<u>EQ 200MG BASE</u>	<u>A202152 004</u>	Mar 27, 2012
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A202152 005</u>	Mar 27, 2012
<u>AB</u>		<u>EQ 400MG BASE</u>	<u>A202152 006</u>	Mar 27, 2012
<u>AB</u>	ALKEM LABS LTD	<u>EQ 25MG BASE</u>	<u>A201504 001</u>	Feb 12, 2013
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A201504 002</u>	Feb 12, 2013
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A201504 003</u>	Feb 12, 2013
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A201504 004</u>	Feb 12, 2013
<u>AB</u>		<u>EQ 200MG BASE</u>	<u>A201504 005</u>	Feb 12, 2013
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A201504 006</u>	Feb 12, 2013
<u>AB</u>		<u>EQ 400MG BASE</u>	<u>A201504 007</u>	Feb 12, 2013
<u>AB</u>	AUROBINDO PHARMA LTD	<u>EQ 25MG BASE</u>	<u>A091388 001</u>	Mar 27, 2012
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A091388 002</u>	Mar 27, 2012
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A091388 003</u>	Mar 27, 2012
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A091388 004</u>	Mar 27, 2012
<u>AB</u>		<u>EQ 200MG BASE</u>	<u>A091388 005</u>	Mar 27, 2012
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A091388 006</u>	Mar 27, 2012
<u>AB</u>		<u>EQ 400MG BASE</u>	<u>A091388 007</u>	Mar 27, 2012
<u>AB</u>	DR REDDYS LABS LTD	<u>EQ 25MG BASE</u>	<u>A077380 001</u>	Mar 27, 2012
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A077380 002</u>	Mar 27, 2012
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A077380 003</u>	Mar 27, 2012
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A077380 004</u>	Mar 27, 2012
<u>AB</u>		<u>EQ 200MG BASE</u>	<u>A077380 005</u>	Mar 27, 2012
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A077380 006</u>	Mar 27, 2012
<u>AB</u>		<u>EQ 400MG BASE</u>	<u>A077380 007</u>	Mar 27, 2012
<u>AB</u>	HETERO LABS LTD V	<u>EQ 25MG BASE</u>	<u>A204316 001</u>	Jun 16, 2022
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A204316 002</u>	Jun 16, 2022
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A204316 003</u>	Jun 16, 2022
<u>AB</u>		<u>EQ 200MG BASE</u>	<u>A204316 004</u>	Jun 16, 2022
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A204316 005</u>	Jun 16, 2022
<u>AB</u>		<u>EQ 400MG BASE</u>	<u>A204316 006</u>	Jun 16, 2022
<u>AB</u>	HIKMA	<u>EQ 25MG BASE</u>	<u>A090120 001</u>	Mar 27, 2012
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A090749 001</u>	Mar 27, 2012
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A090749 002</u>	Mar 27, 2012
<u>AB</u>		<u>EQ 200MG BASE</u>	<u>A090749 003</u>	Mar 27, 2012
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A090749 004</u>	Mar 27, 2012
<u>AB</u>		<u>EQ 400MG BASE</u>	<u>A090749 005</u>	Mar 27, 2012
<u>AB</u>	LUPIN LTD	<u>EQ 25MG BASE</u>	<u>A201109 001</u>	Mar 27, 2012
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A201109 002</u>	Mar 27, 2012
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A201109 003</u>	Mar 27, 2012
<u>AB</u>		<u>EQ 200MG BASE</u>	<u>A201109 004</u>	Mar 27, 2012
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A201109 005</u>	Mar 27, 2012
<u>AB</u>		<u>EQ 400MG BASE</u>	<u>A201109 006</u>	Mar 27, 2012
<u>AB</u>	MACLEODS PHARMS LTD	<u>EQ 25MG BASE</u>	<u>A203359 001</u>	May 17, 2016
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A203359 002</u>	May 17, 2016
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A203359 003</u>	May 17, 2016
<u>AB</u>		<u>EQ 200MG BASE</u>	<u>A203359 004</u>	May 17, 2016
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A203359 005</u>	May 17, 2016
<u>AB</u>		<u>EQ 400MG BASE</u>	<u>A203359 006</u>	May 17, 2016
<u>AB</u>	PRINSTON INC	<u>EQ 25MG BASE</u>	<u>A206954 001</u>	Aug 24, 2022
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A206954 002</u>	Aug 24, 2022
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A206954 003</u>	Aug 24, 2022
<u>AB</u>		<u>EQ 200MG BASE</u>	<u>A206954 004</u>	Aug 24, 2022
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A206954 005</u>	Aug 24, 2022
<u>AB</u>		<u>EQ 400MG BASE</u>	<u>A206954 006</u>	Aug 24, 2022
<u>AB</u>	SCIEGEN PHARMS INC	<u>EQ 25MG BASE</u>	<u>A090960 001</u>	Mar 27, 2012
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A090960 002</u>	Mar 27, 2012
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A090960 003</u>	Mar 27, 2012
<u>AB</u>		<u>EQ 200MG BASE</u>	<u>A090960 004</u>	Mar 27, 2012
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A090960 005</u>	Mar 27, 2012
<u>AB</u>		<u>EQ 400MG BASE</u>	<u>A090960 006</u>	Mar 27, 2012
<u>AB</u>	SUN PHARM	<u>EQ 25MG BASE</u>	<u>A201190 001</u>	Mar 27, 2012

PRESCRIPTION DRUG PRODUCT LIST

QUETIAPINE FUMARATE

TABLET; ORAL

QUETIAPINE FUMARATE

<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A201190 002</u>	Mar 27, 2012	
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A201190 003</u>	Mar 27, 2012	
<u>AB</u>		<u>EQ 200MG BASE</u>	<u>A201190 004</u>	Mar 27, 2012	
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A201190 005</u>	Mar 27, 2012	
<u>AB</u>		<u>EQ 400MG BASE</u>	<u>A201190 006</u>	Mar 27, 2012	
<u>AB</u>	TEVA PHARMS	<u>EQ 25MG BASE</u>	<u>A077745 001</u>	Mar 27, 2012	
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A077745 002</u>	Mar 27, 2012	
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A077745 003</u>	Mar 27, 2012	
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A077745 004</u>	Mar 27, 2012	
<u>AB</u>		<u>EQ 200MG BASE</u>	<u>A077745 005</u>	Mar 27, 2012	
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A077745 006</u>	Mar 27, 2012	
<u>AB</u>		<u>EQ 400MG BASE</u>	<u>A077745 007</u>	Mar 27, 2012	
<u>AB</u>	UNICHEM	<u>EQ 25MG BASE</u>	<u>A202674 001</u>	Mar 08, 2016	
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A202674 002</u>	Mar 08, 2016	
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A202674 003</u>	Mar 08, 2016	
<u>AB</u>		<u>EQ 200MG BASE</u>	<u>A202674 004</u>	Mar 08, 2016	
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A202674 005</u>	Mar 08, 2016	
<u>AB</u>		<u>EQ 400MG BASE</u>	<u>A202674 006</u>	Mar 08, 2016	
<u>SEROQUEL</u>					
<u>AB</u>	+!	<u>ASTRAZENECA</u>	<u>EQ 25MG BASE</u>	<u>N020639 001</u>	Sep 26, 1997
<u>AB</u>	+		<u>EQ 50MG BASE</u>	<u>N020639 007</u>	Oct 04, 2005
<u>AB</u>	+		<u>EQ 100MG BASE</u>	<u>N020639 002</u>	Sep 26, 1997
<u>AB</u>	+		<u>EQ 200MG BASE</u>	<u>N020639 003</u>	Sep 26, 1997
<u>AB</u>	+!		<u>EQ 300MG BASE</u>	<u>N020639 005</u>	Jul 26, 2000
<u>AB</u>	+		<u>EQ 400MG BASE</u>	<u>N020639 006</u>	Oct 04, 2005

TABLET, EXTENDED RELEASE; ORAL

QUETIAPINE FUMARATE

<u>AB</u>	ACCORD HLTHCARE	<u>EQ 50MG BASE</u>	<u>A206252 001</u>	Nov 29, 2017
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A090681 001</u>	May 09, 2017
<u>AB</u>		<u>EQ 200MG BASE</u>	<u>A090681 002</u>	May 09, 2017
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A090681 003</u>	May 09, 2017
<u>AB</u>		<u>EQ 400MG BASE</u>	<u>A090681 004</u>	Nov 01, 2016
<u>AB</u>	ALIGNSCIENCE PHARMA	<u>EQ 150MG BASE</u>	<u>A209497 001</u>	Sep 28, 2018
<u>AB</u>		<u>EQ 200MG BASE</u>	<u>A209497 002</u>	Sep 28, 2018
<u>AB</u>	AUROBINDO PHARMA	<u>EQ 50MG BASE</u>	<u>A207655 001</u>	Nov 29, 2017
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A207655 002</u>	Nov 29, 2017
<u>AB</u>		<u>EQ 200MG BASE</u>	<u>A207655 003</u>	Nov 29, 2017
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A207655 004</u>	Nov 29, 2017
<u>AB</u>		<u>EQ 400MG BASE</u>	<u>A207655 005</u>	Nov 29, 2017
<u>AB</u>	GRAVITI PHARMS	<u>EQ 50MG BASE</u>	<u>A211144 001</u>	Nov 26, 2024
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A211144 002</u>	Nov 26, 2024
<u>AB</u>		<u>EQ 200MG BASE</u>	<u>A211144 003</u>	Nov 26, 2024
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A211144 004</u>	Nov 26, 2024
<u>AB</u>		<u>EQ 400MG BASE</u>	<u>A211144 005</u>	Nov 26, 2024
<u>AB</u>	INTELLIPHARMACEUTIC S	<u>EQ 50MG BASE</u>	<u>A202939 001</u>	May 09, 2017
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A202939 002</u>	May 09, 2017
<u>AB</u>		<u>EQ 200MG BASE</u>	<u>A202939 003</u>	May 09, 2017
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A202939 004</u>	May 09, 2017
<u>AB</u>		<u>EQ 400MG BASE</u>	<u>A202939 005</u>	May 09, 2017
<u>AB</u>	LUPIN LTD	<u>EQ 50MG BASE</u>	<u>A204203 001</u>	May 17, 2017
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A204203 002</u>	May 17, 2017
<u>AB</u>		<u>EQ 200MG BASE</u>	<u>A204203 003</u>	May 17, 2017
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A204203 004</u>	May 17, 2017
<u>AB</u>		<u>EQ 400MG BASE</u>	<u>A204203 005</u>	May 17, 2017
<u>AB</u>	MACLEODS PHARMS LTD	<u>EQ 150MG BASE</u>	<u>A204253 001</u>	Nov 29, 2017
<u>AB</u>		<u>EQ 200MG BASE</u>	<u>A204253 002</u>	Nov 29, 2017
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A204253 003</u>	Nov 29, 2017
<u>AB</u>		<u>EQ 400MG BASE</u>	<u>A204253 004</u>	Nov 29, 2017
<u>AB</u>	NOVAST LABS	<u>EQ 50MG BASE</u>	<u>A208947 001</u>	Nov 29, 2017
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A208947 002</u>	Nov 29, 2017
<u>AB</u>		<u>EQ 200MG BASE</u>	<u>A208947 003</u>	Nov 29, 2017
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A208947 004</u>	Nov 29, 2017
<u>AB</u>		<u>EQ 400MG BASE</u>	<u>A208947 005</u>	Nov 29, 2017
<u>AB</u>	PRINSTON INC	<u>EQ 50MG BASE</u>	<u>A208781 001</u>	Apr 26, 2022
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A208781 002</u>	Apr 26, 2022
<u>AB</u>		<u>EQ 200MG BASE</u>	<u>A208781 003</u>	Apr 26, 2022
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A208781 004</u>	Apr 26, 2022
<u>AB</u>		<u>EQ 400MG BASE</u>	<u>A208781 005</u>	Mar 11, 2024
<u>AB</u>	SCIEGEN PHARMS INC	<u>EQ 50MG BASE</u>	<u>A209635 005</u>	Nov 16, 2018

PRESCRIPTION DRUG PRODUCT LIST

QUETIAPINE FUMARATE

TABLET, EXTENDED RELEASE;ORAL

QUETIAPINE FUMARATE

<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A209635 001</u>	Nov 29, 2017
<u>AB</u>		<u>EQ 200MG BASE</u>	<u>A209635 002</u>	Nov 29, 2017
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A209635 003</u>	Nov 29, 2017
<u>AB</u>		<u>EQ 400MG BASE</u>	<u>A209635 004</u>	Nov 29, 2017
<u>AB</u>	UNICHEM	<u>EQ 50MG BASE</u>	<u>A215478 001</u>	Aug 15, 2022
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A215478 002</u>	Aug 15, 2022
<u>AB</u>		<u>EQ 200MG BASE</u>	<u>A215478 003</u>	Aug 15, 2022
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A215478 004</u>	Aug 15, 2022
<u>AB</u>		<u>EQ 400MG BASE</u>	<u>A215478 005</u>	Aug 15, 2022

SEROQUEL XR

<u>AB</u>	+	ASTRAZENECA	<u>EQ 50MG BASE</u>	<u>N022047 001</u>	May 17, 2007
<u>AB</u>	+		<u>EQ 150MG BASE</u>	<u>N022047 005</u>	Aug 11, 2008
<u>AB</u>	+	!	<u>EQ 200MG BASE</u>	<u>N022047 002</u>	May 17, 2007
<u>AB</u>	+		<u>EQ 300MG BASE</u>	<u>N022047 003</u>	May 17, 2007
<u>AB</u>	+		<u>EQ 400MG BASE</u>	<u>N022047 004</u>	May 17, 2007

QUINAPRIL HYDROCHLORIDE

TABLET;ORAL

QUINAPRIL HYDROCHLORIDE

<u>AB</u>		AUROBINDO PHARMA LTD	<u>EQ 5MG BASE</u>	<u>A202725 001</u>	Apr 29, 2013
<u>AB</u>			<u>EQ 10MG BASE</u>	<u>A202725 002</u>	Apr 29, 2013
<u>AB</u>			<u>EQ 20MG BASE</u>	<u>A202725 003</u>	Apr 29, 2013
<u>AB</u>			<u>EQ 40MG BASE</u>	<u>A202725 004</u>	Apr 29, 2013
<u>AB</u>		CHARTWELL RX	<u>EQ 5MG BASE</u>	<u>A076803 001</u>	Mar 02, 2005
<u>AB</u>			<u>EQ 10MG BASE</u>	<u>A076803 002</u>	Mar 02, 2005
<u>AB</u>			<u>EQ 20MG BASE</u>	<u>A076803 003</u>	Mar 02, 2005
<u>AB</u>			<u>EQ 40MG BASE</u>	<u>A076803 004</u>	Mar 02, 2005
<u>AB</u>		LUPIN	<u>EQ 5MG BASE</u>	<u>A077690 001</u>	Jun 20, 2006
<u>AB</u>			<u>EQ 10MG BASE</u>	<u>A077690 002</u>	Jun 20, 2006
<u>AB</u>			<u>EQ 20MG BASE</u>	<u>A077690 003</u>	Jun 20, 2006
<u>AB</u>	!		<u>EQ 40MG BASE</u>	<u>A077690 004</u>	Jun 20, 2006
<u>AB</u>		PRINSTON INC	<u>EQ 5MG BASE</u>	<u>A205823 001</u>	Sep 15, 2016
<u>AB</u>			<u>EQ 10MG BASE</u>	<u>A205823 002</u>	Sep 15, 2016
<u>AB</u>			<u>EQ 20MG BASE</u>	<u>A205823 003</u>	Sep 15, 2016
<u>AB</u>			<u>EQ 40MG BASE</u>	<u>A205823 004</u>	Sep 15, 2016

QUINIDINE GLUCONATE

TABLET, EXTENDED RELEASE;ORAL

QUINIDINE GLUCONATE

<u>AB</u>		HIBROW HLTHCARE	<u>324MG</u>	<u>A212589 001</u>	Sep 17, 2021
<u>AB</u>	!	SUN PHARM INDUSTRIES	<u>324MG</u>	<u>A089338 001</u>	Feb 11, 1987

QUINIDINE SULFATE

TABLET;ORAL

QUINIDINE SULFATE

!	EPIC PHARMA LLC	200MG	A088072 002	
!		300MG	A088072 001	Sep 26, 1983

QUININE SULFATE

CAPSULE;ORAL

QUININE SULFATE

<u>AB</u>		AMNEAL PHARMS	<u>324MG</u>	<u>A203729 001</u>	Jul 15, 2015
<u>AB</u>		LUPIN LTD	<u>324MG</u>	<u>A203112 001</u>	Apr 24, 2015
<u>AB</u>	!	NOVAST LABS	<u>324MG</u>	<u>A204372 001</u>	Jul 22, 2015
<u>AB</u>		TEVA PHARMS	<u>324MG</u>	<u>A091661 001</u>	Sep 28, 2012

QUIZARTINIB DIHYDROCHLORIDE

TABLET;ORAL

VANFLYTA

+	DAIICHI SANKYO INC	EQ 17.7MG BASE	N216993 001	Jul 20, 2023
+	!	EQ 26.5MG BASE	N216993 002	Jul 20, 2023

RABEPRAZOLE SODIUM

TABLET, DELAYED RELEASE;ORAL

ACIPHEX

<u>AB</u>	+	WAYLIS THERAP	<u>20MG</u>	<u>N020973 002</u>	Aug 19, 1999
-----------	---	---------------	-------------	--------------------	--------------

RABEPRAZOLE SODIUM

<u>AB</u>		ALKEM LABS LTD	<u>20MG</u>	<u>A208644 001</u>	Apr 24, 2018
<u>AB</u>		AMNEAL PHARMS	<u>20MG</u>	<u>A204179 001</u>	Jul 31, 2015
<u>AB</u>		AUROBINDO PHARMA	<u>20MG</u>	<u>A205761 001</u>	Feb 17, 2017
<u>AB</u>		CHARTWELL RX	<u>20MG</u>	<u>A078964 001</u>	Nov 08, 2013

PRESCRIPTION DRUG PRODUCT LIST

RABEPRAZOLE SODIUM

TABLET, DELAYED RELEASE;ORAL

RABEPRAZOLE SODIUM

AB	DR REDDYS	20MG	A076824 001	Nov 08, 2013
AB	LANNETT CO INC	20MG	A090678 001	Nov 08, 2013
AB	RUBICON	20MG	A204237 001	Nov 18, 2015
AB	TORRENT	20MG	A202376 001	Nov 08, 2013

RADIUM RA-223 DICHLORIDE

SOLUTION;INTRAVENOUS

XOFIGO

+	BAYER HLTHCARE	162mCi/6ML (27mCi/ML)	N203971 001	May 15, 2013
---	----------------	-----------------------	-------------	--------------

RALOXIFENE HYDROCHLORIDE

TABLET;ORAL

EVISTA

AB	+	LILLY	60MG	N020815 001	Dec 09, 1997
-----------	---	-------	-------------	--------------------	--------------

RALOXIFENE HYDROCHLORIDE

AB	AMNEAL PHARMS	60MG	A208206 001	Apr 08, 2016
AB	AUROBINDO PHARMA	60MG	A204310 001	Aug 28, 2015
AB	CADILA PHARMS LTD	60MG	A211324 001	Aug 18, 2020
AB	GLENMARK PHARMS LTD	60MG	A204491 001	Mar 22, 2016
AB	INVAGEN PHARMS	60MG	A090842 001	Sep 24, 2014
AB	SCIEGEN PHARMS INC	60MG	A206384 001	Oct 12, 2016
AB	TEVA PHARMS USA	60MG	A078193 001	Mar 04, 2014
AB	WATSON LABS INC	60MG	A200825 001	Jan 21, 2015

RALTEGRAVIR POTASSIUM

POWDER;ORAL

ISENTRESS

+	MSD SUB MERCK	EQ 100MG BASE/PACKET	N205786 001	Dec 20, 2013
---	---------------	----------------------	-------------	--------------

TABLET;ORAL

ISENTRESS

AB	+	MSD SUB MERCK	EQ 400MG BASE	N022145 001	Oct 12, 2007
-----------	---	---------------	----------------------	--------------------	--------------

RALTEGRAVIR POTASSIUM

AB	HETERO LABS LTD III	EQ 400MG BASE	A203540 001	Dec 19, 2024
	ISENTRESS HD			
+	MSD SUB MERCK	EQ 600MG BASE	N022145 002	May 26, 2017
	TABLET, CHEWABLE;ORAL			
	ISENTRESS			
+	MSD SUB MERCK	EQ 25MG BASE	N203045 001	Dec 21, 2011
+		EQ 100MG BASE	N203045 002	Dec 21, 2011

RAMELTEON

TABLET;ORAL

RAMELTEON

AB	ACTAVIS LABS FL INC	8MG	A091610 001	Aug 19, 2015
AB	ANDAS 5 HOLDING	8MG	A215435 001	Aug 24, 2022
AB	AUROBINDO PHARMA LTD	8MG	A215972 001	Jul 10, 2023
AB	DR REDDYS LABS SA	8MG	A091693 001	Jul 26, 2013
AB	GRANULES	8MG	A213186 001	Aug 21, 2020
AB	I3 PHARMS	8MG	A212650 001	Apr 10, 2020
AB	MICRO LABS	8MG	A215243 001	Feb 09, 2023
AB	UPSHER SMITH LABS	8MG	A213815 001	Oct 26, 2020
AB	XIROMED	8MG	A216209 001	Nov 25, 2022
AB	ZYDUS PHARMS	8MG	A211567 001	Jul 22, 2019

ROZEREM

AB	+	TAKEDA PHARMS USA	8MG	N021782 001	Jul 22, 2005
-----------	---	-------------------	------------	--------------------	--------------

RAMIPRIL

CAPSULE;ORAL

ALTACE

AB	+	KING PHARMS LLC	1.25MG	N019901 001	Jan 28, 1991
AB	+		2.5MG	N019901 002	Jan 28, 1991
AB	+		5MG	N019901 003	Jan 28, 1991
AB	+		10MG	N019901 004	Jan 28, 1991

RAMIPRIL

AB	AUROBINDO PHARMA LTD	1.25MG	A091604 001	Jun 08, 2011
AB		2.5MG	A091604 002	Jun 08, 2011
AB		5MG	A091604 003	Jun 08, 2011
AB		10MG	A091604 004	Jun 08, 2011
AB	CHARTWELL MOLECULAR	1.25MG	A078745 001	Jun 18, 2008
AB		2.5MG	A078745 002	Jun 18, 2008

PRESCRIPTION DRUG PRODUCT LIST

RAMIPRIL

CAPSULE;ORAL

RAMIPRIL

<u>AB</u>		<u>5MG</u>	<u>A078745 003</u>	Jun 18, 2008
<u>AB</u>		<u>10MG</u>	<u>A078745 004</u>	Jun 18, 2008
<u>AB</u>	COREPHARMA	<u>1.25MG</u>	<u>A079116 001</u>	Jun 20, 2008
<u>AB</u>		<u>2.5MG</u>	<u>A079116 002</u>	Jun 20, 2008
<u>AB</u>		<u>5MG</u>	<u>A079116 003</u>	Jun 20, 2008
<u>AB</u>		<u>10MG</u>	<u>A079116 004</u>	Jun 20, 2008
<u>AB</u>	DR REDDYS LABS LTD	<u>1.25MG</u>	<u>A078191 001</u>	Jun 18, 2008
<u>AB</u>		<u>2.5MG</u>	<u>A078191 002</u>	Jun 18, 2008
<u>AB</u>		<u>5MG</u>	<u>A078191 003</u>	Jun 18, 2008
<u>AB</u>		<u>10MG</u>	<u>A078191 004</u>	Jun 18, 2008
<u>AB</u>	HIKMA	<u>1.25MG</u>	<u>A077900 001</u>	Jun 18, 2008
<u>AB</u>		<u>2.5MG</u>	<u>A077900 002</u>	Jun 18, 2008
<u>AB</u>		<u>5MG</u>	<u>A077900 003</u>	Jun 18, 2008
<u>AB</u>		<u>10MG</u>	<u>A077900 004</u>	Jun 18, 2008
<u>AB</u>	LUPIN	<u>1.25MG</u>	<u>A077626 001</u>	Jun 09, 2008
<u>AB</u>		<u>2.5MG</u>	<u>A077626 002</u>	Jun 09, 2008
<u>AB</u>		<u>5MG</u>	<u>A077626 003</u>	Jun 09, 2008
<u>AB</u>		<u>10MG</u>	<u>A077626 004</u>	Jun 09, 2008
<u>AB</u>	WATSON LABS	<u>1.25MG</u>	<u>A076549 001</u>	Oct 24, 2005
<u>AB</u>		<u>2.5MG</u>	<u>A076549 002</u>	Oct 24, 2005
<u>AB</u>		<u>10MG</u>	<u>A076549 004</u>	Oct 24, 2005
<u>AB</u>	ZYDUS PHARMS USA	<u>1.25MG</u>	<u>A078832 001</u>	Sep 02, 2008
<u>AB</u>		<u>2.5MG</u>	<u>A078832 002</u>	Sep 02, 2008
<u>AB</u>		<u>5MG</u>	<u>A078832 003</u>	Sep 02, 2008
<u>AB</u>		<u>10MG</u>	<u>A078832 004</u>	Sep 02, 2008

RANITIDINE HYDROCHLORIDE

CAPSULE;ORAL

RANITIDINE HYDROCHLORIDE

<u>AB</u>	DR REDDYS LABS LTD	<u>EQ 150MG BASE</u>	<u>A075742 001</u>	Nov 29, 2000
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A075742 002</u>	Nov 29, 2000
<u>AB</u>	SANDOZ	<u>EQ 150MG BASE</u>	<u>A074655 001</u>	Oct 22, 1997
<u>AB</u>	!	<u>EQ 300MG BASE</u>	<u>A074655 002</u>	Oct 22, 1997

TABLET;ORAL

RANITIDINE HYDROCHLORIDE

<u>AB</u>	APOTEX	<u>EQ 150MG BASE</u>	<u>A074680 001</u>	Sep 12, 1997
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A074680 002</u>	Sep 12, 1997
<u>AB</u>	DR REDDYS LABS INC	<u>EQ 150MG BASE</u>	<u>A076705 001</u>	Jul 27, 2005
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A076705 002</u>	Jul 27, 2005
<u>AB</u>	GLENMARK PHARMS INC	<u>EQ 150MG BASE</u>	<u>A078542 001</u>	Nov 19, 2008
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A078542 002</u>	Nov 19, 2008
<u>AB</u>	SANDOZ	<u>EQ 150MG BASE</u>	<u>A074467 001</u>	Aug 29, 1997
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A074467 002</u>	Aug 29, 1997
<u>AB</u>	VKT PHARMA	<u>EQ 150MG BASE</u>	<u>A211289 001</u>	Jan 31, 2019
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A211289 002</u>	Jan 31, 2019

RANOLAZINE

TABLET, EXTENDED RELEASE;ORAL

RANOLAZINE

<u>AB</u>	ACTAVIS ELIZABETH	<u>500MG</u>	<u>A208862 001</u>	May 28, 2019
<u>AB</u>		<u>1GM</u>	<u>A208862 002</u>	May 28, 2019
<u>AB</u>	AJANTA PHARMA LTD	<u>500MG</u>	<u>A210054 001</u>	May 28, 2019
<u>AB</u>		<u>1GM</u>	<u>A210054 002</u>	May 28, 2019
<u>AB</u>	ALKEM LABS LTD	<u>500MG</u>	<u>A209953 001</u>	Nov 30, 2020
<u>AB</u>		<u>1GM</u>	<u>A209953 002</u>	Nov 30, 2020
<u>AB</u>	AUROBINDO PHARMA	<u>500MG</u>	<u>A209081 001</u>	Dec 23, 2022
<u>AB</u>		<u>1GM</u>	<u>A209081 002</u>	Dec 23, 2022
<u>AB</u>	CADILA	<u>500MG</u>	<u>A210188 001</u>	Aug 19, 2019
<u>AB</u>		<u>1GM</u>	<u>A210188 002</u>	Aug 19, 2019
<u>AB</u>	CHARTWELL RX	<u>500MG</u>	<u>A201046 001</u>	Jul 29, 2013
<u>AB</u>		<u>1GM</u>	<u>A201046 002</u>	Jul 29, 2013
<u>AB</u>	GLENMARK PHARMS LTD	<u>500MG</u>	<u>A211082 001</u>	Jul 05, 2019
<u>AB</u>		<u>1GM</u>	<u>A211082 002</u>	Jul 05, 2019
<u>AB</u>	HETERO LABS LTD III	<u>500MG</u>	<u>A212788 001</u>	May 05, 2022
<u>AB</u>		<u>1GM</u>	<u>A212788 002</u>	May 05, 2022
<u>AB</u>	I3 PHARMS	<u>500MG</u>	<u>A213517 001</u>	Apr 27, 2022
<u>AB</u>		<u>1GM</u>	<u>A213517 002</u>	Apr 27, 2022
<u>AB</u>	MANKIND PHARMA	<u>500MG</u>	<u>A212284 001</u>	Feb 12, 2020
<u>AB</u>		<u>1GM</u>	<u>A212284 002</u>	Feb 12, 2020
<u>AB</u>	MICRO LABS	<u>500MG</u>	<u>A211745 001</u>	Feb 27, 2020

PRESCRIPTION DRUG PRODUCT LIST

RANOLAZINE

TABLET, EXTENDED RELEASE;ORAL

RANOLAZINE

<u>AB</u>		<u>1GM</u>	<u>A211745</u>	<u>002</u>	Feb 27, 2020
<u>AB</u>	NOVAST LABS	<u>500MG</u>	<u>A210668</u>	<u>001</u>	Sep 27, 2023
<u>AB</u>		<u>1GM</u>	<u>A210668</u>	<u>002</u>	Sep 27, 2023
<u>AB</u>	PRAXGEN	<u>1GM</u>	<u>A212781</u>	<u>002</u>	Mar 23, 2020
<u>AB</u>		<u>500MG</u>	<u>A212781</u>	<u>001</u>	Mar 23, 2020
<u>AB</u>	RISING	<u>500MG</u>	<u>A212889</u>	<u>001</u>	Jan 28, 2021
<u>AB</u>		<u>1GM</u>	<u>A212889</u>	<u>002</u>	Jan 28, 2021
<u>AB</u>	SCIEGEN PHARMS INC	<u>500MG</u>	<u>A211829</u>	<u>001</u>	Jun 04, 2019
<u>AB</u>		<u>1GM</u>	<u>A211829</u>	<u>002</u>	Jun 04, 2019
<u>AB</u>	SUN PHARM	<u>500MG</u>	<u>A211707</u>	<u>001</u>	May 28, 2019
<u>AB</u>	!	<u>1GM</u>	<u>A211707</u>	<u>002</u>	May 28, 2019
<u>AB</u>	SUNSHINE	<u>500MG</u>	<u>A211865</u>	<u>001</u>	Mar 23, 2020
<u>AB</u>		<u>1GM</u>	<u>A211865</u>	<u>002</u>	Mar 23, 2020
<u>AB</u>	UNICHEM	<u>500MG</u>	<u>A213083</u>	<u>001</u>	Mar 16, 2023
<u>AB</u>		<u>1GM</u>	<u>A213083</u>	<u>002</u>	Mar 16, 2023
<u>AB</u>	VKT PHARMA	<u>500MG</u>	<u>A214035</u>	<u>001</u>	Jan 19, 2022
<u>AB</u>		<u>1GM</u>	<u>A214035</u>	<u>002</u>	Jan 19, 2022

RASAGILINE MESYLATE

TABLET;ORAL

AZILECT

<u>AB</u>	+	TEVA	<u>EQ 0.5MG BASE</u>	<u>N021641</u>	<u>001</u>	May 16, 2006
<u>AB</u>	+	!	<u>EQ 1MG BASE</u>	<u>N021641</u>	<u>002</u>	May 16, 2006

RASAGILINE MESYLATE

<u>AB</u>		ALKEM LABS LTD	<u>EQ 0.5MG BASE</u>	<u>A201889</u>	<u>001</u>	Oct 30, 2017
<u>AB</u>			<u>EQ 1MG BASE</u>	<u>A201889</u>	<u>002</u>	Oct 30, 2017
<u>AB</u>		AUROBINDO PHARMA USA	<u>EQ 0.5MG BASE</u>	<u>A201971</u>	<u>001</u>	May 15, 2017
<u>AB</u>			<u>EQ 1MG BASE</u>	<u>A201971</u>	<u>002</u>	May 15, 2017
<u>AB</u>		CHARTWELL RX	<u>EQ 0.5MG BASE</u>	<u>A201892</u>	<u>001</u>	Jul 27, 2018
<u>AB</u>			<u>EQ 1MG BASE</u>	<u>A201892</u>	<u>002</u>	Jul 27, 2018
<u>AB</u>		INDOCO	<u>EQ 0.5MG BASE</u>	<u>A206153</u>	<u>001</u>	Oct 04, 2019
<u>AB</u>			<u>EQ 1MG BASE</u>	<u>A206153</u>	<u>002</u>	Oct 04, 2019
<u>AB</u>		MACLEODS PHARMS LTD	<u>EQ 0.5MG BASE</u>	<u>A208866</u>	<u>001</u>	Jun 28, 2024
<u>AB</u>			<u>EQ 1MG BASE</u>	<u>A208866</u>	<u>002</u>	Jun 28, 2024
<u>AB</u>		MICRO LABS	<u>EQ 0.5MG BASE</u>	<u>A207004</u>	<u>001</u>	Mar 29, 2019
<u>AB</u>			<u>EQ 1MG BASE</u>	<u>A207004</u>	<u>002</u>	Mar 29, 2019
<u>AB</u>		ORBION PHARMS	<u>EQ 0.5MG BASE</u>	<u>A201970</u>	<u>001</u>	Mar 15, 2016
<u>AB</u>			<u>EQ 1MG BASE</u>	<u>A201970</u>	<u>002</u>	Mar 15, 2016
<u>AB</u>		SKG PHARMA	<u>EQ 0.5MG BASE</u>	<u>A218163</u>	<u>001</u>	Apr 09, 2024
<u>AB</u>			<u>EQ 1MG BASE</u>	<u>A218163</u>	<u>002</u>	Apr 09, 2024

REGADENOSON

SOLUTION;INTRAVENOUS

LEXISCAN

<u>AP</u>	+	!	ASTELLAS	<u>0.4MG/5ML (0.08MG/ML)</u>	<u>N022161</u>	<u>001</u>	Apr 10, 2008
-----------	---	---	----------	------------------------------	----------------	------------	--------------

REGADENOSON

<u>AP</u>			ACCORD HLTHCARE	<u>0.4MG/5ML (0.08MG/ML)</u>	<u>A213236</u>	<u>001</u>	Apr 11, 2022
<u>AP</u>			APOTEX	<u>0.4MG/5ML (0.08MG/ML)</u>	<u>A207604</u>	<u>001</u>	Apr 11, 2022
<u>AP</u>			BAXTER HLTHCARE CORP	<u>0.4MG/5ML (0.08MG/ML)</u>	<u>A217455</u>	<u>001</u>	May 23, 2023
<u>AP</u>			DR REDDYS	<u>0.4MG/5ML (0.08MG/ML)</u>	<u>A213210</u>	<u>001</u>	Apr 11, 2022
<u>AP</u>			EUGIA PHARMA	<u>0.4MG/5ML (0.08MG/ML)</u>	<u>A216437</u>	<u>001</u>	Oct 26, 2022
<u>AP</u>			GE HEALTHCARE	<u>0.4MG/5ML (0.08MG/ML)</u>	<u>A215955</u>	<u>001</u>	Aug 05, 2024
<u>AP</u>			GLAND PHARMA LTD	<u>0.4MG/5ML (0.08MG/ML)</u>	<u>A207320</u>	<u>001</u>	Jul 12, 2022
<u>AP</u>			HIKMA	<u>0.4MG/5ML (0.08MG/ML)</u>	<u>A215827</u>	<u>001</u>	Feb 02, 2023
<u>AP</u>			HOSPIRA	<u>0.4MG/5ML (0.08MG/ML)</u>	<u>A214349</u>	<u>001</u>	Aug 31, 2022
<u>AP</u>			IMS LTD	<u>0.4MG/5ML (0.08MG/ML)</u>	<u>A214252</u>	<u>001</u>	May 23, 2022
<u>AP</u>			INDIES PHARMA	<u>0.4MG/5ML (0.08MG/ML)</u>	<u>A218054</u>	<u>001</u>	Aug 22, 2024
<u>AP</u>			MEITHEAL	<u>0.4MG/5ML (0.08MG/ML)</u>	<u>A212806</u>	<u>001</u>	Apr 11, 2022
<u>AP</u>			MYLAN	<u>0.4MG/5ML (0.08MG/ML)</u>	<u>A213856</u>	<u>001</u>	Apr 04, 2023

REGORAFENIB

TABLET;ORAL

STIVARGA

	+	!	BAYER HLTHCARE	40MG	N203085	001	Sep 27, 2012
--	---	---	----------------	------	---------	-----	--------------

PRESCRIPTION DRUG PRODUCT LIST

RELUGOLIX

TABLET; ORAL

ORGOVYX

+! SUMITOMO PHARMA 120MG N214621 001 Dec 18, 2020

REMDESIVIR

POWDER; INTRAVENOUS

VEKLURY

+! GILEAD SCIENCES INC 100MG/VIAL N214787 001 Oct 22, 2020

SOLUTION; INTRAVENOUS

VEKLURY

+! GILEAD SCIENCES INC 100MG/20ML (5MG/ML) N214787 002 Oct 22, 2020

REMIFENTANIL HYDROCHLORIDE

INJECTABLE; INJECTION

REMIFENTANIL HYDROCHLORIDEAP FRESENIUS KABI USA EQ 1MG BASE/VIAL A206223 001 Jan 16, 2018AP EQ 2MG BASE/VIAL A206223 002 Jan 16, 2018AP EQ 5MG BASE/VIAL A206223 003 Jan 16, 2018AP HIKMA EQ 1MG BASE/VIAL A210594 001 Oct 13, 2020AP EQ 2MG BASE/VIAL A210594 002 Oct 13, 2020AP EQ 5MG BASE/VIAL A210594 003 Oct 13, 2020AP NIVAGEN PHARMS INC EQ 1MG BASE/VIAL A215635 001 Jun 28, 2024AP EQ 2MG BASE/VIAL A215635 002 Jun 28, 2024AP EQ 5MG BASE/VIAL A215635 003 Jun 28, 2024ULTIVAAP + MYLAN INSTITUTIONAL EQ 1MG BASE/VIAL N020630 001 Jul 12, 1996AP + EQ 2MG BASE/VIAL N020630 002 Jul 12, 1996AP +! EQ 5MG BASE/VIAL N020630 003 Jul 12, 1996REMIMAZOLAM BESYLATE

POWDER; INTRAVENOUS

BYFAVO

+! ACACIA EQ 20MG BASE/VIAL N212295 001 Oct 06, 2020

REPAGLINIDE

TABLET; ORAL

REPAGLINIDEAB AUROBINDO PHARMA LTD 0.5MG A203820 001 Jan 22, 2014AB 1MG A203820 002 Jan 22, 2014AB ! 2MG A203820 003 Jan 22, 2014AB CHARTWELL RX 0.5MG A078555 001 Nov 22, 2013AB 1MG A078555 002 Jan 22, 2014AB 2MG A078555 003 Jan 22, 2014AB MACLEODS PHARMS LTD 0.5MG A207209 001 Mar 22, 2023AB 1MG A207209 002 Mar 22, 2023AB 2MG A207209 003 Mar 22, 2023AB PADAGIS US 0.5MG A201189 001 Jul 17, 2013AB 1MG A201189 002 Jan 22, 2014AB 2MG A201189 003 Jan 22, 2014AB SUN PHARM INDS INC 1MG A077571 002 Jul 11, 2013AB 2MG A077571 003 Jul 11, 2013REPOTRECTINIB

CAPSULE; ORAL

AUGTYRO

+! BRISTOL 40MG N218213 001 Nov 15, 2023

+ 160MG N218213 002 Jun 11, 2024

RESMETIROM

TABLET; ORAL

REZDIFFRA

+ MADRIGAL 60MG N217785 001 Mar 14, 2024

+ 80MG N217785 002 Mar 14, 2024

+! 100MG N217785 003 Mar 14, 2024

REVEFENACIN

SOLUTION; INHALATION

YUPELRI

+! MYLAN IRELAND LTD 175MCG/3ML N210598 001 Nov 09, 2018

PRESCRIPTION DRUG PRODUCT LISTREVUMENIB CITRATE

TABLET; ORAL

REVUFORJ

+	SYNDAX	EQ 25MG BASE	N218944 001	Nov 15, 2024
+		EQ 110MG BASE	N218944 002	Nov 15, 2024
+	!	EQ 160MG BASE	N218944 003	Nov 15, 2024

REZAFUNGIN ACETATE

POWDER; INTRAVENOUS

REZZAYO

+	MUNDIPHARMA	EQ 200MG BASE/VIAL	N217417 001	Mar 22, 2023
---	-------------	--------------------	-------------	--------------

RIBAVIRIN

CAPSULE; ORAL

RIBAVIRIN

<u>AB</u>	!	AUROBINDO PHARMA	<u>200MG</u>	<u>A079117 001</u>	Sep 17, 2009
<u>AB</u>		ZYDUS PHARMS USA	<u>200MG</u>	<u>A077224 001</u>	Oct 28, 2005

FOR SOLUTION; INHALATION

RIBAVIRIN

<u>AN</u>		NAVINTA LLC	<u>6GM/VIAL</u>	<u>A207366 001</u>	Oct 06, 2016
-----------	--	-------------	-----------------	--------------------	--------------

VIRAZOLE

<u>AN</u>	+	BAUSCH	<u>6GM/VIAL</u>	<u>N018859 001</u>	Dec 31, 1985
-----------	---	--------	-----------------	--------------------	--------------

TABLET; ORAL

RIBAVIRIN

<u>AB</u>		AUROBINDO PHARMA	<u>200MG</u>	<u>A079111 001</u>	Sep 17, 2009
<u>AB</u>		SANDOZ	<u>200MG</u>	<u>A077743 001</u>	Oct 03, 2006
<u>AB</u>		ZYDUS PHARMS USA	<u>200MG</u>	<u>A077094 001</u>	Dec 05, 2005

RIBOCICLIB SUCCINATE

TABLET; ORAL

KISQALI

+	NOVARTIS	EQ 200MG BASE	N209092 001	Mar 13, 2017
---	----------	---------------	-------------	--------------

RIBOFLAVIN 5'-PHOSPHATE

SOLUTION/DROPS; OPHTHALMIC

PHOTREXA

+	GLAUKOS	0.146%	N203324 001	Apr 15, 2016
---	---------	--------	-------------	--------------

PHOTREXA VISCOUS IN DEXTRAN 20%

+	GLAUKOS	0.146%	N203324 002	Apr 15, 2016
---	---------	--------	-------------	--------------

RIFABUTIN

CAPSULE; ORAL

MYCOBUTIN

<u>AB</u>	+	PFIZER	<u>150MG</u>	<u>N050689 001</u>	Dec 23, 1992
-----------	---	--------	--------------	--------------------	--------------

RIFABUTIN

<u>AB</u>		LUPIN LTD	<u>150MG</u>	<u>A090033 001</u>	Feb 24, 2014
-----------	--	-----------	--------------	--------------------	--------------

<u>AB</u>		NOVITIUM PHARMA	<u>150MG</u>	<u>A215041 001</u>	Dec 17, 2021
-----------	--	-----------------	--------------	--------------------	--------------

RIFAMPIN

CAPSULE; ORAL

RIFAMPIN

<u>AB</u>		CHARTWELL MOLECULAR	<u>150MG</u>	<u>A065390 001</u>	Mar 28, 2008
-----------	--	---------------------	--------------	--------------------	--------------

<u>AB</u>			<u>300MG</u>	<u>A065390 002</u>	Mar 28, 2008
-----------	--	--	--------------	--------------------	--------------

<u>AB</u>		EPIC PHARMA LLC	<u>150MG</u>	<u>A064150 002</u>	Jan 02, 1998
-----------	--	-----------------	--------------	--------------------	--------------

<u>AB</u>			<u>300MG</u>	<u>A064150 001</u>	May 28, 1997
-----------	--	--	--------------	--------------------	--------------

<u>AB</u>		LUPIN PHARMS	<u>150MG</u>	<u>A090034 001</u>	Aug 21, 2013
-----------	--	--------------	--------------	--------------------	--------------

<u>AB</u>	!		<u>300MG</u>	<u>A090034 002</u>	Aug 21, 2013
-----------	---	--	--------------	--------------------	--------------

RIMACTANE

<u>AB</u>		OXFORD PHARMS	<u>300MG</u>	<u>N050429 001</u>	
-----------	--	---------------	--------------	--------------------	--

INJECTABLE; INJECTION

RIFADIN

<u>AP</u>	+	SANOFI AVENTIS US	<u>600MG/VIAL</u>	<u>N050627 001</u>	May 25, 1989
-----------	---	-------------------	-------------------	--------------------	--------------

RIFAMPIN

<u>AP</u>		EPIC PHARMA LLC	<u>600MG/VIAL</u>	<u>A065502 001</u>	Sep 21, 2010
-----------	--	-----------------	-------------------	--------------------	--------------

<u>AP</u>		FRESENIUS KABI USA	<u>600MG/VIAL</u>	<u>A091181 001</u>	Aug 21, 2014
-----------	--	--------------------	-------------------	--------------------	--------------

<u>AP</u>		HIKMA	<u>600MG/VIAL</u>	<u>A064217 001</u>	Oct 29, 1999
-----------	--	-------	-------------------	--------------------	--------------

<u>AP</u>		HIKMA PHARMS	<u>600MG/VIAL</u>	<u>A205039 001</u>	Mar 03, 2016
-----------	--	--------------	-------------------	--------------------	--------------

<u>AP</u>		MYLAN LABS LTD	<u>600MG/VIAL</u>	<u>A065421 001</u>	May 22, 2008
-----------	--	----------------	-------------------	--------------------	--------------

RIFAPENTINE

TABLET; ORAL

PRIFTIN

+	SANOFI AVENTIS US	150MG	N021024 001	Jun 22, 1998
---	-------------------	-------	-------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

RIFAXIMIN

TABLET;ORAL

XIFAXAN

+	!	SALIX PHARMS	200MG	N021361	001	May 25, 2004
+	!		550MG	N021361	002	Mar 24, 2010

RILPIVIRINE HYDROCHLORIDE

TABLET;ORAL

EDURANT

+	!	JANSSEN PRODS	EQ 25MG BASE	N202022	001	May 20, 2011
---	---	---------------	--------------	---------	-----	--------------

RILUZOLE

SUSPENSION;ORAL

RILUZOLE

AB		ALKEM LABS LTD	50MG/10ML	A216035	001	Aug 22, 2024
-----------	--	----------------	------------------	----------------	------------	--------------

TIGLUTIK KIT

AB	+	!	ITALFARMACO SA	50MG/10ML	N209080	001	Sep 05, 2018
-----------	---	---	----------------	------------------	----------------	------------	--------------

TABLET;ORAL

RILUZOLE

AB	!	ALKEM LABS LTD	50MG	A204048	001	Mar 30, 2016
-----------	---	----------------	-------------	----------------	------------	--------------

AB		GLENMARK PHARMS LTD	50MG	A091394	001	Jun 18, 2013
-----------	--	---------------------	-------------	----------------	------------	--------------

AB		IMPAX LABS	50MG	A076173	001	Jan 29, 2003
-----------	--	------------	-------------	----------------	------------	--------------

AB		KENTON	50MG	A206045	001	Dec 09, 2019
-----------	--	--------	-------------	----------------	------------	--------------

AB		MYLAN PHARMS INC	50MG	A203042	001	Jul 01, 2013
-----------	--	------------------	-------------	----------------	------------	--------------

RIMANTADINE HYDROCHLORIDE

TABLET;ORAL

RIMANTADINE HYDROCHLORIDE

!	!	IMPAX LABS	100MG	A076132	001	Aug 30, 2002
---	---	------------	-------	---------	-----	--------------

RIMEGEPANT SULFATE

TABLET, ORALLY DISINTEGRATING;ORAL

NURTEC ODT

+	!	PFIZER	EQ 75MG BASE	N212728	001	Feb 27, 2020
---	---	--------	--------------	---------	-----	--------------

RIOCIGUAT

TABLET;ORAL

ADEMPAS

AB	+		0.5MG	N204819	001	Oct 08, 2013
-----------	---	--	--------------	----------------	------------	--------------

AB	+		1MG	N204819	002	Oct 08, 2013
-----------	---	--	------------	----------------	------------	--------------

AB	+		1.5MG	N204819	003	Oct 08, 2013
-----------	---	--	--------------	----------------	------------	--------------

AB	+		2MG	N204819	004	Oct 08, 2013
-----------	---	--	------------	----------------	------------	--------------

AB	+	!	2.5MG	N204819	005	Oct 08, 2013
-----------	---	---	--------------	----------------	------------	--------------

RIOCIGUAT

AB		MSN	0.5MG	A211135	001	Sep 01, 2022
-----------	--	-----	--------------	----------------	------------	--------------

AB			1MG	A211135	002	Sep 01, 2022
-----------	--	--	------------	----------------	------------	--------------

AB			1.5MG	A211135	003	Sep 01, 2022
-----------	--	--	--------------	----------------	------------	--------------

AB			2MG	A211135	004	Sep 01, 2022
-----------	--	--	------------	----------------	------------	--------------

AB			2.5MG	A211135	005	Sep 01, 2022
-----------	--	--	--------------	----------------	------------	--------------

RIPRETINIB

TABLET;ORAL

QINLOCK

+	!	DECIPHERA PHARMS	50MG	N213973	001	May 15, 2020
---	---	------------------	------	---------	-----	--------------

RISDIPLAM

FOR SOLUTION;ORAL

EVRYSDI

+	!	GENENTECH INC	0.75MG/ML	N213535	001	Aug 07, 2020
---	---	---------------	-----------	---------	-----	--------------

RISEDRONATE SODIUM

TABLET;ORAL

ACTONEL

AB	+	!	APIL	35MG	N020835	003	May 25, 2002
-----------	---	---	------	-------------	----------------	------------	--------------

AB	+	!		150MG	N020835	005	Apr 22, 2008
-----------	---	---	--	--------------	----------------	------------	--------------

RISEDRONATE SODIUM

AB		APOTEX	35MG	A090877	001	Nov 30, 2015
-----------	--	--------	-------------	----------------	------------	--------------

AB			75MG	A090877	002	Jun 10, 2014
-----------	--	--	-------------	----------------	------------	--------------

AB			150MG	A090877	003	Jun 10, 2014
-----------	--	--	--------------	----------------	------------	--------------

AB		AUROBINDO PHARMA LTD	5MG	A200296	001	Nov 30, 2015
-----------	--	----------------------	------------	----------------	------------	--------------

AB			30MG	A200296	002	Nov 30, 2015
-----------	--	--	-------------	----------------	------------	--------------

AB			35MG	A200296	003	Nov 30, 2015
-----------	--	--	-------------	----------------	------------	--------------

AB		MACLEODS PHARMS LTD	5MG	A203533	001	Dec 09, 2015
-----------	--	---------------------	------------	----------------	------------	--------------

AB			30MG	A203533	002	Dec 09, 2015
-----------	--	--	-------------	----------------	------------	--------------

AB			35MG	A203533	003	Nov 29, 2016
-----------	--	--	-------------	----------------	------------	--------------

AB		ORBION PHARMS	30MG	A205280	001	May 13, 2019
-----------	--	---------------	-------------	----------------	------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

RISEDRONATE SODIUM

TABLET; ORAL

RISEDRONATE SODIUM

<u>AB</u>		<u>35MG</u>	<u>A205280</u>	<u>002</u>	May 13, 2019
<u>AB</u>	SUN PHARM	<u>5MG</u>	<u>A090886</u>	<u>001</u>	Nov 30, 2015
<u>AB</u>		<u>30MG</u>	<u>A090886</u>	<u>002</u>	Nov 30, 2015
<u>AB</u>		<u>35MG</u>	<u>A090886</u>	<u>003</u>	Nov 30, 2015
<u>AB</u>		<u>75MG</u>	<u>A090886</u>	<u>004</u>	Jun 10, 2014
<u>AB</u>		<u>150MG</u>	<u>A090886</u>	<u>005</u>	Jun 10, 2014
<u>AB</u>	TEVA PHARMS USA	<u>5MG</u>	<u>A077132</u>	<u>001</u>	Oct 05, 2007
<u>AB</u>		<u>30MG</u>	<u>A077132</u>	<u>002</u>	Oct 05, 2007
<u>AB</u>		<u>35MG</u>	<u>A077132</u>	<u>003</u>	Oct 05, 2007
<u>AB</u>		<u>150MG</u>	<u>A079215</u>	<u>001</u>	Jun 13, 2014
BX	AUROBINDO PHARMA	150MG	A206768	001	Oct 21, 2016

TABLET, DELAYED RELEASE; ORAL

ATELVIA

<u>AB</u>	+!	APIL	<u>35MG</u>	<u>N022560</u>	<u>001</u>	Oct 08, 2010
-----------	----	------	-------------	----------------	------------	--------------

RISEDRONATE SODIUM

<u>AB</u>		SUN PHARM	<u>35MG</u>	<u>A203925</u>	<u>001</u>	Jul 09, 2019
<u>AB</u>		TEVA PHARMS USA	<u>35MG</u>	<u>A203217</u>	<u>001</u>	May 18, 2015

RISPERIDONE

FOR SUSPENSION, EXTENDED RELEASE; INTRAMUSCULAR

RYKINDO

+	SHANDONG LUYE	12.5MG	N212849	001	Jan 13, 2023
+!		25MG	N212849	002	Jan 13, 2023
+		37.5MG	N212849	003	Jan 13, 2023
+		50MG	N212849	004	Jan 13, 2023

FOR SUSPENSION, EXTENDED RELEASE; SUBCUTANEOUS

PERSERIS KIT

+	INDIVIOR	90MG	N210655	001	Jul 27, 2018
+!		120MG	N210655	002	Jul 27, 2018

INJECTABLE; INTRAMUSCULAR

RISPERDAL CONSTA

<u>AB</u>	+	JANSSEN PHARMS	<u>12.5MG/VIAL</u>	<u>N021346</u>	<u>004</u>	Apr 12, 2007
<u>AB</u>	+!		<u>25MG/VIAL</u>	<u>N021346</u>	<u>001</u>	Oct 29, 2003
<u>AB</u>	+		<u>37.5MG/VIAL</u>	<u>N021346</u>	<u>002</u>	Oct 29, 2003
<u>AB</u>	+		<u>50MG/VIAL</u>	<u>N021346</u>	<u>003</u>	Oct 29, 2003

RISPERIDONE

<u>AB</u>		TEVA PHARMS USA INC	<u>12.5MG/VIAL</u>	<u>A214068</u>	<u>001</u>	Dec 05, 2023
<u>AB</u>			<u>25MG/VIAL</u>	<u>A214068</u>	<u>002</u>	Dec 05, 2023
<u>AB</u>			<u>37.5MG/VIAL</u>	<u>A214068</u>	<u>003</u>	Dec 05, 2023
<u>AB</u>			<u>50MG/VIAL</u>	<u>A214068</u>	<u>004</u>	Dec 05, 2023

SOLUTION; ORAL

RISPERDAL

<u>AA</u>	+!	JANSSEN PHARMS	<u>1MG/ML</u>	<u>N020588</u>	<u>001</u>	Jun 10, 1996
-----------	----	----------------	---------------	----------------	------------	--------------

RISPERIDONE

<u>AA</u>		AMNEAL PHARMS	<u>1MG/ML</u>	<u>A091384</u>	<u>001</u>	May 25, 2011
<u>AA</u>		AUROBINDO PHARMA LTD	<u>1MG/ML</u>	<u>A078452</u>	<u>001</u>	Sep 04, 2009
<u>AA</u>		CHARTWELL MOLECULAR	<u>1MG/ML</u>	<u>A202386</u>	<u>001</u>	Jan 12, 2015
<u>AA</u>		LANNETT CO INC	<u>1MG/ML</u>	<u>A079158</u>	<u>001</u>	Dec 03, 2010
<u>AA</u>		TARO	<u>1MG/ML</u>	<u>A090347</u>	<u>001</u>	Feb 07, 2011
<u>AA</u>		TRIS PHARMA INC	<u>1MG/ML</u>	<u>A079059</u>	<u>001</u>	Dec 12, 2012

SUSPENSION, EXTENDED RELEASE; SUBCUTANEOUS

UZEDY

+	TEVA	50MG/0.14ML (50MG/0.14ML)	N213586	001	Apr 28, 2023
+		75MG/0.21ML (75MG/0.21ML)	N213586	002	Apr 28, 2023
+		100MG/0.28ML (100MG/0.28ML)	N213586	003	Apr 28, 2023
+		125MG/0.35ML (125MG/0.35ML)	N213586	004	Apr 28, 2023
+		150MG/0.42ML (150MG/0.42ML)	N213586	005	Apr 28, 2023
+		200MG/0.56ML (200MG/0.56ML)	N213586	006	Apr 28, 2023
+		250MG/0.7ML (250MG/0.7ML)	N213586	007	Apr 28, 2023

TABLET; ORAL

RISPERDAL

<u>AB</u>	+	JANSSEN PHARMS	<u>0.25MG</u>	<u>N020272</u>	<u>008</u>	May 10, 1999
<u>AB</u>	+		<u>0.5MG</u>	<u>N020272</u>	<u>007</u>	Jan 27, 1999
<u>AB</u>	+!		<u>1MG</u>	<u>N020272</u>	<u>001</u>	Dec 29, 1993
<u>AB</u>	+		<u>2MG</u>	<u>N020272</u>	<u>002</u>	Dec 29, 1993
<u>AB</u>	+		<u>3MG</u>	<u>N020272</u>	<u>003</u>	Dec 29, 1993
<u>AB</u>	+		<u>4MG</u>	<u>N020272</u>	<u>004</u>	Dec 29, 1993

RISPERIDONE

<u>AB</u>		AJANTA PHARMA LTD	<u>0.25MG</u>	<u>A201003</u>	<u>001</u>	Aug 24, 2011
-----------	--	-------------------	---------------	----------------	------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

RISPERIDONE

TABLET; ORAL

RISPERIDONE

<u>AB</u>		<u>0.5MG</u>	<u>A201003 002</u>	Aug 24, 2011
<u>AB</u>		<u>1MG</u>	<u>A201003 003</u>	Aug 24, 2011
<u>AB</u>		<u>2MG</u>	<u>A201003 004</u>	Aug 24, 2011
<u>AB</u>		<u>3MG</u>	<u>A201003 005</u>	Aug 24, 2011
<u>AB</u>		<u>4MG</u>	<u>A201003 006</u>	Aug 24, 2011
<u>AB</u>	AMNEAL	<u>0.25MG</u>	<u>A078071 001</u>	Jun 17, 2009
<u>AB</u>		<u>0.5MG</u>	<u>A078071 002</u>	Jun 17, 2009
<u>AB</u>		<u>1MG</u>	<u>A078071 003</u>	Jun 17, 2009
<u>AB</u>		<u>2MG</u>	<u>A078071 004</u>	Jun 17, 2009
<u>AB</u>		<u>3MG</u>	<u>A078071 005</u>	Jun 17, 2009
<u>AB</u>		<u>4MG</u>	<u>A078071 006</u>	Jun 17, 2009
<u>AB</u>	APOTEX INC	<u>0.25MG</u>	<u>A077953 001</u>	Sep 15, 2008
<u>AB</u>		<u>0.5MG</u>	<u>A077953 002</u>	Sep 15, 2008
<u>AB</u>		<u>1MG</u>	<u>A077953 003</u>	Sep 15, 2008
<u>AB</u>		<u>2MG</u>	<u>A077953 004</u>	Sep 15, 2008
<u>AB</u>		<u>3MG</u>	<u>A077953 005</u>	Sep 15, 2008
<u>AB</u>		<u>4MG</u>	<u>A077953 006</u>	Sep 15, 2008
<u>AB</u>	CHARTWELL MOLECULAR	<u>0.25MG</u>	<u>A077543 001</u>	May 18, 2011
<u>AB</u>		<u>0.5MG</u>	<u>A077543 002</u>	May 18, 2011
<u>AB</u>		<u>1MG</u>	<u>A077543 003</u>	May 18, 2011
<u>AB</u>		<u>2MG</u>	<u>A077543 004</u>	May 18, 2011
<u>AB</u>		<u>3MG</u>	<u>A077543 005</u>	May 18, 2011
<u>AB</u>		<u>4MG</u>	<u>A077543 006</u>	May 18, 2011
<u>AB</u>	DR REDDYS LABS LTD	<u>0.25MG</u>	<u>A076879 001</u>	Oct 24, 2008
<u>AB</u>		<u>0.5MG</u>	<u>A076879 002</u>	Oct 24, 2008
<u>AB</u>		<u>1MG</u>	<u>A076879 003</u>	Oct 24, 2008
<u>AB</u>		<u>2MG</u>	<u>A076879 004</u>	Oct 24, 2008
<u>AB</u>		<u>3MG</u>	<u>A076879 005</u>	Oct 24, 2008
<u>AB</u>		<u>4MG</u>	<u>A076879 006</u>	Oct 24, 2008
<u>AB</u>	ESJAY PHARMA	<u>0.25MG</u>	<u>A078871 001</u>	Oct 09, 2008
<u>AB</u>		<u>0.5MG</u>	<u>A078871 002</u>	Oct 09, 2008
<u>AB</u>		<u>1MG</u>	<u>A078871 003</u>	Oct 09, 2008
<u>AB</u>		<u>2MG</u>	<u>A078871 004</u>	Oct 09, 2008
<u>AB</u>		<u>3MG</u>	<u>A078871 005</u>	Oct 09, 2008
<u>AB</u>		<u>4MG</u>	<u>A078871 006</u>	Oct 09, 2008
<u>AB</u>	IPCA LABS LTD	<u>0.25MG</u>	<u>A205104 001</u>	Jun 26, 2024
<u>AB</u>		<u>0.5MG</u>	<u>A205104 002</u>	Jun 26, 2024
<u>AB</u>		<u>1MG</u>	<u>A205104 003</u>	Jun 26, 2024
<u>AB</u>		<u>2MG</u>	<u>A205104 004</u>	Jun 26, 2024
<u>AB</u>		<u>3MG</u>	<u>A205104 005</u>	Jun 26, 2024
<u>AB</u>		<u>4MG</u>	<u>A205104 006</u>	Jun 26, 2024
<u>AB</u>	PRINSTON INC	<u>0.25MG</u>	<u>A077493 001</u>	Nov 29, 2011
<u>AB</u>		<u>0.5MG</u>	<u>A077493 002</u>	Nov 29, 2011
<u>AB</u>		<u>1MG</u>	<u>A077493 003</u>	Nov 29, 2011
<u>AB</u>		<u>2MG</u>	<u>A077493 004</u>	Nov 29, 2011
<u>AB</u>		<u>3MG</u>	<u>A077493 005</u>	Nov 29, 2011
<u>AB</u>		<u>4MG</u>	<u>A077493 006</u>	Nov 29, 2011
<u>AB</u>	RENATA	<u>0.25MG</u>	<u>A078707 001</u>	Dec 29, 2008
<u>AB</u>		<u>0.5MG</u>	<u>A078707 002</u>	Dec 29, 2008
<u>AB</u>		<u>1MG</u>	<u>A078707 003</u>	Dec 29, 2008
<u>AB</u>		<u>2MG</u>	<u>A078707 004</u>	Dec 29, 2008
<u>AB</u>		<u>3MG</u>	<u>A078707 005</u>	Dec 29, 2008
<u>AB</u>		<u>4MG</u>	<u>A078707 006</u>	Dec 29, 2008
<u>AB</u>	RISING	<u>0.25MG</u>	<u>A078269 001</u>	Oct 08, 2008
<u>AB</u>		<u>0.5MG</u>	<u>A078269 002</u>	Oct 08, 2008
<u>AB</u>		<u>1MG</u>	<u>A078269 003</u>	Oct 08, 2008
<u>AB</u>		<u>2MG</u>	<u>A078269 004</u>	Oct 08, 2008
<u>AB</u>		<u>3MG</u>	<u>A078269 005</u>	Oct 08, 2008
<u>AB</u>		<u>4MG</u>	<u>A078269 006</u>	Oct 08, 2008
<u>AB</u>	SANDOZ	<u>0.25MG</u>	<u>A078528 001</u>	Oct 16, 2009
<u>AB</u>		<u>0.5MG</u>	<u>A078528 002</u>	Oct 16, 2009
<u>AB</u>		<u>1MG</u>	<u>A078528 003</u>	Oct 16, 2009
<u>AB</u>		<u>2MG</u>	<u>A078528 004</u>	Oct 16, 2009
<u>AB</u>		<u>3MG</u>	<u>A078528 005</u>	Oct 16, 2009
<u>AB</u>		<u>4MG</u>	<u>A078528 006</u>	Oct 16, 2009
<u>AB</u>	TORRENT PHARMS	<u>0.25MG</u>	<u>A079088 001</u>	Oct 30, 2008
<u>AB</u>		<u>0.5MG</u>	<u>A079088 002</u>	Oct 30, 2008
<u>AB</u>		<u>1MG</u>	<u>A079088 003</u>	Oct 30, 2008
<u>AB</u>		<u>2MG</u>	<u>A079088 004</u>	Oct 30, 2008

PRESCRIPTION DRUG PRODUCT LIST

RISPERIDONE

TABLET; ORAL

RISPERIDONE

<u>AB</u>		<u>3MG</u>	<u>A079088</u>	<u>005</u>	Oct 30, 2008
<u>AB</u>		<u>4MG</u>	<u>A079088</u>	<u>006</u>	Oct 30, 2008
<u>AB</u>	ZYDUS PHARMS USA INC	<u>0.25MG</u>	<u>A078040</u>	<u>001</u>	Oct 16, 2008
<u>AB</u>		<u>0.5MG</u>	<u>A078040</u>	<u>002</u>	Oct 16, 2008
<u>AB</u>		<u>1MG</u>	<u>A078040</u>	<u>003</u>	Oct 16, 2008
<u>AB</u>		<u>2MG</u>	<u>A078040</u>	<u>004</u>	Oct 16, 2008
<u>AB</u>		<u>3MG</u>	<u>A078040</u>	<u>005</u>	Oct 16, 2008
<u>AB</u>		<u>4MG</u>	<u>A078040</u>	<u>006</u>	Oct 16, 2008

TABLET, ORALLY DISINTEGRATING; ORAL

RISPERIDONE

<u>AB</u>	DR REDDYS LABS LTD	<u>0.5MG</u>	<u>A077328</u>	<u>001</u>	Feb 24, 2009
<u>AB</u>		<u>1MG</u>	<u>A077328</u>	<u>002</u>	Oct 05, 2009
<u>AB</u>		<u>2MG</u>	<u>A077328</u>	<u>003</u>	Feb 24, 2009
<u>AB</u>		<u>3MG</u>	<u>A077328</u>	<u>004</u>	Nov 30, 2009
<u>AB</u>		<u>4MG</u>	<u>A077328</u>	<u>005</u>	Nov 30, 2009
<u>AB</u>	ENDO OPERATIONS	<u>0.5MG</u>	<u>A077494</u>	<u>002</u>	Apr 30, 2009
<u>AB</u>		<u>1MG</u>	<u>A077494</u>	<u>003</u>	Oct 26, 2009
<u>AB</u>		<u>2MG</u>	<u>A077494</u>	<u>004</u>	Apr 30, 2009
<u>AB</u>		<u>3MG</u>	<u>A077494</u>	<u>005</u>	Apr 30, 2009
<u>AB</u>		<u>4MG</u>	<u>A077494</u>	<u>006</u>	Apr 30, 2009
<u>AB</u>	JUBILANT GENERICS	<u>0.5MG</u>	<u>A090839</u>	<u>001</u>	Nov 04, 2011
<u>AB</u>	!	<u>1MG</u>	<u>A090839</u>	<u>002</u>	Nov 04, 2011
<u>AB</u>		<u>2MG</u>	<u>A090839</u>	<u>003</u>	Nov 04, 2011
<u>AB</u>		<u>3MG</u>	<u>A090839</u>	<u>004</u>	Nov 04, 2011
<u>AB</u>		<u>4MG</u>	<u>A090839</u>	<u>005</u>	Nov 04, 2011
<u>AB</u>	SANDOZ	<u>0.5MG</u>	<u>A078116</u>	<u>001</u>	Dec 22, 2009
<u>AB</u>		<u>1MG</u>	<u>A078116</u>	<u>002</u>	Dec 22, 2009
<u>AB</u>		<u>2MG</u>	<u>A078116</u>	<u>003</u>	Dec 22, 2009
<u>AB</u>		<u>3MG</u>	<u>A078116</u>	<u>004</u>	Dec 22, 2009
<u>AB</u>		<u>4MG</u>	<u>A078116</u>	<u>005</u>	Dec 22, 2009
<u>AB</u>	SUN PHARM INDS LTD	<u>0.5MG</u>	<u>A077542</u>	<u>001</u>	Aug 06, 2010
<u>AB</u>		<u>0.5MG</u>	<u>A078464</u>	<u>001</u>	Apr 08, 2013
<u>AB</u>		<u>1MG</u>	<u>A077542</u>	<u>002</u>	Aug 06, 2010
<u>AB</u>		<u>1MG</u>	<u>A078464</u>	<u>002</u>	Apr 08, 2013
<u>AB</u>		<u>2MG</u>	<u>A077542</u>	<u>003</u>	Aug 06, 2010
<u>AB</u>		<u>2MG</u>	<u>A078464</u>	<u>003</u>	Apr 08, 2013
<u>AB</u>		<u>3MG</u>	<u>A078464</u>	<u>004</u>	Apr 08, 2013
<u>AB</u>		<u>3MG</u>	<u>A078474</u>	<u>001</u>	Aug 06, 2010
<u>AB</u>		<u>4MG</u>	<u>A078464</u>	<u>005</u>	Apr 08, 2013
<u>AB</u>		<u>4MG</u>	<u>A078474</u>	<u>002</u>	Aug 06, 2010
<u>AB</u>	ZYDUS PHARMS USA	<u>0.5MG</u>	<u>A078516</u>	<u>001</u>	May 01, 2009
<u>AB</u>		<u>2MG</u>	<u>A078516</u>	<u>003</u>	May 01, 2009
<u>AB</u>	ENDO OPERATIONS	0.25MG	A077494	001	Apr 30, 2009

RITLECITINIB TOSYLATE

CAPSULE; ORAL

LITFULO

+! PFIZER

EQ 50MG BASE

N215830 001 Jun 23, 2023

RITONAVIR

CAPSULE; ORAL

RITONAVIR

NORVIUM BIOSCIENCE

100MG

A205024 001 Feb 16, 2024

POWDER; ORAL

NORVIR

+! ABBVIE

100MG/PACKET

N209512 001 Jun 07, 2017

TABLET; ORAL

NORVIR

<u>AB</u>	+! ABBVIE	<u>100MG</u>	<u>N022417</u>	<u>001</u>	Feb 10, 2010
-----------	-----------	--------------	----------------	------------	--------------

RITONAVIR

<u>AB</u>	AMNEAL	<u>100MG</u>	<u>A208890</u>	<u>001</u>	Sep 17, 2018
<u>AB</u>	AUROBINDO PHARMA LTD	<u>100MG</u>	<u>A206614</u>	<u>001</u>	Sep 17, 2018
<u>AB</u>	CIPLA	<u>100MG</u>	<u>A202573</u>	<u>001</u>	Jan 15, 2015
<u>AB</u>		<u>100MG</u>	<u>A203759</u>	<u>001</u>	May 10, 2022
<u>AB</u>	HETERO LABS LTD III	<u>100MG</u>	<u>A204587</u>	<u>001</u>	Sep 17, 2018

PRESCRIPTION DRUG PRODUCT LIST

RIVAROXABAN

FOR SUSPENSION;ORAL

XARELTO

+! JANSSEN PHARMS 1MG/ML N215859 001 Dec 20, 2021

TABLET;ORAL

XARELTO

+ JANSSEN PHARMS 2.5MG N022406 004 Oct 11, 2018

+ 10MG N022406 001 Jul 01, 2011

+ 15MG N022406 002 Nov 04, 2011

+! 20MG N022406 003 Nov 04, 2011

RIVASTIGMINE

FILM, EXTENDED RELEASE;TRANSDERMAL

EXELONAB + SANDOZ 4.6MG/24HR N022083 001 Jul 06, 2007AB +! 9.5MG/24HR N022083 002 Jul 06, 2007AB + 13.3MG/24HR N022083 005 Aug 31, 2012RIVASTIGMINEAB ALVOGEN 4.6MG/24HR A204403 001 Sep 03, 2015AB 9.5MG/24HR A204403 002 Sep 03, 2015AB 13.3MG/24HR A204403 003 Aug 31, 2015AB AMNEAL PHARMS 4.6MG/24HR A207308 001 Jan 08, 2019AB 9.5MG/24HR A207308 002 Jan 08, 2019AB 13.3MG/24HR A207308 003 Jan 08, 2019AB BRECKENRIDGE 4.6MG/24HR A209063 001 Nov 26, 2019AB 9.5MG/24HR A209063 002 Nov 26, 2019AB 13.3MG/24HR A209063 003 Nov 26, 2019AB MYLAN TECHNOLOGIES 4.6MG/24HR A205622 001 Jun 20, 2018AB 9.5MG/24HR A205622 002 Jun 20, 2018AB 13.3MG/24HR A205622 003 Jun 20, 2018AB ZYDUS PHARMS 4.6MG/24HR A206318 001 Mar 04, 2019AB 9.5MG/24HR A206318 002 Mar 04, 2019AB 13.3MG/24HR A206318 003 Mar 04, 2019RIVASTIGMINE TARTRATE

CAPSULE;ORAL

RIVASTIGMINE TARTRATEAB ALEMBIC PHARMS LTD EQ 1.5MG BASE A091689 001 Jun 12, 2012AB EQ 3MG BASE A091689 002 Jun 12, 2012AB EQ 4.5MG BASE A091689 003 Jun 12, 2012AB EQ 6MG BASE A091689 004 Jun 12, 2012AB AUROBINDO PHARMA EQ 1.5MG BASE A204572 001 Mar 25, 2016AB EQ 3MG BASE A204572 002 Mar 25, 2016AB EQ 4.5MG BASE A204572 003 Mar 25, 2016AB EQ 6MG BASE A204572 004 Mar 25, 2016AB CADILA PHARMS LTD EQ 1.5MG BASE A203844 001 Feb 13, 2017AB EQ 3MG BASE A203844 002 Feb 13, 2017AB EQ 4.5MG BASE A203844 003 Feb 13, 2017AB EQ 6MG BASE A203844 004 Feb 13, 2017AB CHARTWELL RX EQ 1.5MG BASE A207797 001 Sep 28, 2017AB EQ 3MG BASE A207797 002 Sep 28, 2017AB EQ 4.5MG BASE A207797 003 Sep 28, 2017AB EQ 6MG BASE A207797 004 Sep 28, 2017AB ! DR REDDYS LABS INC EQ 1.5MG BASE A077130 001 Oct 31, 2007AB EQ 3MG BASE A077130 002 Oct 31, 2007AB EQ 4.5MG BASE A077130 003 Oct 31, 2007AB ! EQ 6MG BASE A077130 004 Oct 31, 2007AB MACLEODS PHARMS LTD EQ 1.5MG BASE A203148 001 Aug 22, 2014AB EQ 3MG BASE A203148 002 Aug 22, 2014AB EQ 4.5MG BASE A203148 003 Aug 22, 2014AB EQ 6MG BASE A203148 004 Aug 22, 2014AB ORBION PHARMS EQ 1.5MG BASE A090879 001 Jun 10, 2015AB EQ 3MG BASE A090879 002 Jun 10, 2015AB EQ 4.5MG BASE A090879 003 Jun 10, 2015AB EQ 6MG BASE A090879 004 Jun 10, 2015AB WATSON LABS EQ 1.5MG BASE A077129 001 Jan 08, 2008AB EQ 3MG BASE A077129 002 Jan 08, 2008AB EQ 4.5MG BASE A077129 003 Jan 08, 2008AB EQ 6MG BASE A077129 004 Jan 08, 2008

PRESCRIPTION DRUG PRODUCT LIST

RIZATRIPTAN BENZOATE

TABLET; ORAL

MAXALT

<u>AB</u>	<u>+!</u> ORGANON LLC	<u>EQ 10MG BASE</u>	<u>N020864 002</u>	Jun 29, 1998
<u>RIZATRIPTAN BENZOATE</u>				
<u>AB</u>	ALKEM LABS LTD	<u>EQ 5MG BASE</u>	<u>A203269 001</u>	Feb 18, 2016
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A203269 002</u>	Feb 18, 2016
<u>AB</u>	AUROBINDO PHARMA LTD	<u>EQ 5MG BASE</u>	<u>A202490 001</u>	Dec 31, 2012
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A202490 002</u>	Dec 31, 2012
<u>AB</u>	CHARTWELL RX	<u>EQ 5MG BASE</u>	<u>A077526 001</u>	Mar 26, 2013
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A077526 002</u>	Mar 26, 2013
<u>AB</u>	CREEKWOOD PHARMS	<u>EQ 5MG BASE</u>	<u>A202047 001</u>	Dec 31, 2012
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A202047 002</u>	Dec 31, 2012
<u>AB</u>	MACLEODS PHARMS LTD	<u>EQ 5MG BASE</u>	<u>A203147 001</u>	Feb 11, 2014
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A203147 002</u>	Feb 11, 2014
<u>AB</u>	MYLAN PHARMS INC	<u>EQ 5MG BASE</u>	<u>A201993 001</u>	Dec 31, 2012
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A201993 002</u>	Dec 31, 2012
<u>AB</u>	TEVA PHARMS	<u>EQ 5MG BASE</u>	<u>A077263 001</u>	Dec 31, 2012
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A077263 002</u>	Dec 31, 2012

TABLET, ORALLY DISINTEGRATING; ORAL

MAXALT-MLT

<u>AB</u>	<u>+!</u> ORGANON	<u>EQ 10MG BASE</u>	<u>N020865 002</u>	Jun 29, 1998
<u>RIZATRIPTAN BENZOATE</u>				
<u>AB</u>	AUROBINDO PHARMA LTD	<u>EQ 5MG BASE</u>	<u>A203062 001</u>	Jul 01, 2013
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A203062 002</u>	Jul 01, 2013
<u>AB</u>	MACLEODS PHARMS LTD	<u>EQ 5MG BASE</u>	<u>A203146 001</u>	Sep 19, 2014
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A203146 002</u>	Sep 19, 2014
<u>AB</u>	NATCO PHARMA LTD	<u>EQ 5MG BASE</u>	<u>A203478 001</u>	Jul 01, 2013
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A203478 002</u>	Jul 01, 2013
<u>AB</u>	PANACEA	<u>EQ 5MG BASE</u>	<u>A204722 001</u>	Jan 11, 2017
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A204722 002</u>	Jan 11, 2017

ROCURONIUM BROMIDE

INJECTABLE; INJECTION

ROCURONIUM BROMIDE

<u>AP</u>	CAPLIN	<u>50MG/5ML (10MG/ML)</u>	<u>A216234 001</u>	Mar 02, 2023
<u>AP</u>		<u>100MG/10ML (10MG/ML)</u>	<u>A216234 002</u>	Mar 02, 2023
<u>AP</u>	EUGIA PHARMA	<u>50MG/5ML (10MG/ML)</u>	<u>A206206 001</u>	Apr 12, 2017
<u>AP</u>		<u>100MG/10ML (10MG/ML)</u>	<u>A206206 002</u>	Apr 12, 2017
<u>AP</u>	FRESENIUS KABI USA	<u>50MG/5ML (10MG/ML)</u>	<u>A078651 001</u>	Dec 29, 2008
<u>AP</u>		<u>100MG/10ML (10MG/ML)</u>	<u>A078651 002</u>	Dec 29, 2008
<u>AP</u>	<u>!</u> GLAND PHARMA LTD	<u>50MG/5ML (10MG/ML)</u>	<u>A205656 001</u>	Apr 04, 2018
<u>AP</u>	<u>!</u>	<u>100MG/10ML (10MG/ML)</u>	<u>A205656 002</u>	Apr 04, 2018
<u>AP</u>	HOSPIRA	<u>50MG/5ML (10MG/ML)</u>	<u>A078519 001</u>	Nov 26, 2008
<u>AP</u>		<u>100MG/10ML (10MG/ML)</u>	<u>A078519 002</u>	Nov 26, 2008
<u>AP</u>	MEITHEAL	<u>50MG/5ML (10MG/ML)</u>	<u>A213453 001</u>	May 08, 2023
<u>AP</u>		<u>100MG/10ML (10MG/ML)</u>	<u>A213453 002</u>	May 08, 2023
<u>AP</u>	MYLAN INSTITUTIONAL	<u>50MG/5ML (10MG/ML)</u>	<u>A079199 001</u>	Nov 26, 2008
<u>AP</u>		<u>100MG/10ML (10MG/ML)</u>	<u>A079199 002</u>	Nov 26, 2008
<u>AP</u>	PIRAMAL CRITICAL	<u>50MG/5ML (10MG/ML)</u>	<u>A210437 001</u>	Aug 13, 2019
<u>AP</u>		<u>100MG/10ML (10MG/ML)</u>	<u>A210437 002</u>	Aug 13, 2019
<u>AP</u>	PRINSTON INC	<u>50MG/5ML (10MG/ML)</u>	<u>A212668 001</u>	May 26, 2022
<u>AP</u>		<u>100MG/10ML (10MG/ML)</u>	<u>A212668 002</u>	May 26, 2022
<u>AP</u>	SAGENT PHARMS INC	<u>50MG/5ML (10MG/ML)</u>	<u>A091458 001</u>	Jul 28, 2010
<u>AP</u>		<u>100MG/10ML (10MG/ML)</u>	<u>A091458 002</u>	Jul 28, 2010
<u>AP</u>	SANDOZ	<u>50MG/5ML (10MG/ML)</u>	<u>A079195 001</u>	Dec 05, 2008
<u>AP</u>		<u>100MG/10ML (10MG/ML)</u>	<u>A079195 002</u>	Dec 05, 2008
<u>AP</u>	SHANDONG	<u>50MG/5ML (10MG/ML)</u>	<u>A215684 001</u>	Jul 12, 2024
<u>AP</u>		<u>100MG/10ML (10MG/ML)</u>	<u>A215684 002</u>	Jul 12, 2024
<u>AP</u>	STERISCIENCE	<u>50MG/5ML (10MG/ML)</u>	<u>A217092 001</u>	Jul 22, 2024
<u>AP</u>	TAMARANG	<u>50MG/5ML (10MG/ML)</u>	<u>A091115 001</u>	Aug 27, 2012
<u>AP</u>		<u>100MG/10ML (10MG/ML)</u>	<u>A091115 002</u>	Aug 27, 2012
<u>AP</u>	WEST WARD PHARM CORP	<u>50MG/5ML (10MG/ML)</u>	<u>A204679 001</u>	Feb 28, 2017
<u>AP</u>		<u>100MG/10ML (10MG/ML)</u>	<u>A204679 002</u>	Feb 28, 2017

PRESCRIPTION DRUG PRODUCT LIST

ROPINIROLE HYDROCHLORIDE

TABLET; ORAL

ROPINIROLE HYDROCHLORIDE

<u>AB</u>	MLV	<u>EQ 0.25MG BASE</u>	<u>A079165 001</u>	Feb 07, 2012
<u>AB</u>		<u>EQ 0.5MG BASE</u>	<u>A079165 002</u>	Feb 07, 2012
<u>AB</u>		<u>EQ 1MG BASE</u>	<u>A079165 003</u>	Feb 07, 2012
<u>AB</u>		<u>EQ 2MG BASE</u>	<u>A079165 004</u>	Feb 07, 2012
<u>AB</u>		<u>EQ 3MG BASE</u>	<u>A079165 005</u>	Feb 07, 2012
<u>AB</u>		<u>EQ 4MG BASE</u>	<u>A079165 006</u>	Feb 07, 2012
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A079165 007</u>	Feb 07, 2012
<u>AB</u>	ORBION PHARMS	<u>EQ 0.25MG BASE</u>	<u>A079229 001</u>	Nov 28, 2012
<u>AB</u>		<u>EQ 0.5MG BASE</u>	<u>A079229 002</u>	Nov 28, 2012
<u>AB</u>		<u>EQ 1MG BASE</u>	<u>A079229 003</u>	Nov 28, 2012
<u>AB</u>		<u>EQ 2MG BASE</u>	<u>A079229 004</u>	Nov 28, 2012
<u>AB</u>		<u>EQ 3MG BASE</u>	<u>A079229 005</u>	Nov 28, 2012
<u>AB</u>		<u>EQ 4MG BASE</u>	<u>A079229 006</u>	Nov 28, 2012
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A079229 007</u>	Nov 28, 2012
<u>AB</u>	! PRINSTON INC	<u>EQ 0.25MG BASE</u>	<u>A078110 001</u>	May 05, 2008
<u>AB</u>		<u>EQ 0.5MG BASE</u>	<u>A078110 002</u>	May 05, 2008
<u>AB</u>		<u>EQ 1MG BASE</u>	<u>A078110 003</u>	May 05, 2008
<u>AB</u>		<u>EQ 2MG BASE</u>	<u>A078110 004</u>	May 05, 2008
<u>AB</u>		<u>EQ 3MG BASE</u>	<u>A078110 005</u>	May 05, 2008
<u>AB</u>		<u>EQ 4MG BASE</u>	<u>A078110 006</u>	May 05, 2008
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A078110 007</u>	Jul 11, 2008
<u>AB</u>	ZYDUS LIFESCIENCES	<u>EQ 0.25MG BASE</u>	<u>A090411 001</u>	Jun 01, 2009
<u>AB</u>		<u>EQ 0.5MG BASE</u>	<u>A090411 002</u>	Jun 01, 2009
<u>AB</u>		<u>EQ 1MG BASE</u>	<u>A090411 003</u>	Jun 01, 2009
<u>AB</u>		<u>EQ 2MG BASE</u>	<u>A090411 004</u>	Jun 01, 2009
<u>AB</u>		<u>EQ 3MG BASE</u>	<u>A090411 005</u>	Jun 01, 2009
<u>AB</u>		<u>EQ 4MG BASE</u>	<u>A090411 006</u>	Jun 01, 2009
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A090411 007</u>	Jun 01, 2009

TABLET, EXTENDED RELEASE; ORAL

ROPINIROLE HYDROCHLORIDE

<u>AB</u>	ACTAVIS ELIZABETH	<u>EQ 2MG BASE</u>	<u>A090869 001</u>	May 17, 2012
<u>AB</u>		<u>EQ 4MG BASE</u>	<u>A090869 002</u>	May 17, 2012
<u>AB</u>		<u>EQ 6MG BASE</u>	<u>A090869 003</u>	May 17, 2012
<u>AB</u>		<u>EQ 8MG BASE</u>	<u>A090869 004</u>	May 17, 2012
<u>AB</u>		<u>EQ 12MG BASE</u>	<u>A090869 005</u>	May 17, 2012
<u>AB</u>	ALEMBIC	<u>EQ 2MG BASE</u>	<u>A202786 001</u>	Apr 22, 2013
<u>AB</u>		<u>EQ 4MG BASE</u>	<u>A202786 002</u>	Apr 22, 2013
<u>AB</u>		<u>EQ 6MG BASE</u>	<u>A202786 003</u>	Apr 22, 2013
<u>AB</u>		<u>EQ 8MG BASE</u>	<u>A202786 004</u>	Apr 22, 2013
<u>AB</u>		<u>EQ 12MG BASE</u>	<u>A202786 005</u>	Apr 22, 2013
<u>AB</u>	CHARTWELL RX	<u>EQ 2MG BASE</u>	<u>A091395 001</u>	Aug 27, 2012
<u>AB</u>		<u>EQ 4MG BASE</u>	<u>A091395 002</u>	Aug 27, 2012
<u>AB</u>		<u>EQ 6MG BASE</u>	<u>A091395 003</u>	Aug 27, 2012
<u>AB</u>		<u>EQ 8MG BASE</u>	<u>A091395 004</u>	Aug 27, 2012
<u>AB</u>		<u>EQ 12MG BASE</u>	<u>A091395 005</u>	Aug 27, 2012
<u>AB</u>	! DR REDDYS LABS LTD	<u>EQ 2MG BASE</u>	<u>A201576 001</u>	Jun 06, 2012
<u>AB</u>		<u>EQ 4MG BASE</u>	<u>A201576 002</u>	Jun 06, 2012
<u>AB</u>		<u>EQ 6MG BASE</u>	<u>A201576 003</u>	Jun 06, 2012
<u>AB</u>		<u>EQ 8MG BASE</u>	<u>A201576 004</u>	Jun 06, 2012
<u>AB</u>		<u>EQ 12MG BASE</u>	<u>A201576 005</u>	Jun 06, 2012
<u>AB</u>	PRINSTON INC	<u>EQ 2MG BASE</u>	<u>A204413 001</u>	May 11, 2022
<u>AB</u>		<u>EQ 4MG BASE</u>	<u>A204413 002</u>	May 11, 2022
<u>AB</u>		<u>EQ 6MG BASE</u>	<u>A204413 003</u>	May 11, 2022
<u>AB</u>		<u>EQ 8MG BASE</u>	<u>A204413 004</u>	May 11, 2022
<u>AB</u>		<u>EQ 12MG BASE</u>	<u>A204413 005</u>	May 11, 2022
<u>AB</u>	SANDOZ INC	<u>EQ 2MG BASE</u>	<u>A201047 001</u>	Jun 06, 2012
<u>AB</u>		<u>EQ 4MG BASE</u>	<u>A201047 003</u>	Jun 06, 2012
<u>AB</u>		<u>EQ 6MG BASE</u>	<u>A201047 004</u>	Jun 06, 2012
<u>AB</u>		<u>EQ 8MG BASE</u>	<u>A201047 005</u>	Jun 06, 2012
<u>AB</u>		<u>EQ 12MG BASE</u>	<u>A201047 006</u>	Jun 06, 2012

ROPIVACAINE HYDROCHLORIDE

SOLUTION; INJECTION

NAROPIN

<u>AP</u>	+	FRESENIUS KABI USA	<u>20MG/10ML (2MG/ML)</u>	<u>N020533 001</u>	May 01, 1998
<u>AP</u>	+		<u>40MG/20ML (2MG/ML)</u>	<u>N020533 002</u>	Sep 24, 1996
<u>AP</u>	+		<u>100MG/20ML (5MG/ML)</u>	<u>N020533 003</u>	May 01, 1998
<u>AP</u>	+		<u>100MG/10ML (10MG/ML)</u>	<u>N020533 005</u>	Sep 24, 1996
<u>AP</u>	+		<u>150MG/20ML (7.5MG/ML)</u>	<u>N020533 004</u>	Sep 24, 1996
<u>AP</u>	+		<u>150MG/30ML (5MG/ML)</u>	<u>N020533 008</u>	Sep 24, 1996

PRESCRIPTION DRUG PRODUCT LIST

ROPIVACAINE HYDROCHLORIDE

SOLUTION; INJECTION

NAROPIN

<u>AP</u>	+	<u>200MG/100ML (2MG/ML)</u>	<u>N020533 006</u>	Sep 24, 1996
<u>AP</u>	+	<u>200MG/20ML (10MG/ML)</u>	<u>N020533 011</u>	Sep 24, 1996
<u>AP</u>	+	<u>400MG/200ML (2MG/ML)</u>	<u>N020533 007</u>	Sep 24, 1996
<u>AP</u>	+	<u>500MG/100ML (5MG/ML)</u>	<u>N020533 009</u>	Jan 04, 2011
<u>AP</u>	+	<u>1GM/200ML (5MG/ML)</u>	<u>N020533 010</u>	Jan 04, 2011

ROPIVACAINE HYDROCHLORIDE

<u>AP</u>	AMNEAL	<u>200MG/100ML (2MG/ML)</u>	<u>A216605 001</u>	Mar 08, 2023
<u>AP</u>		<u>400MG/200ML (2MG/ML)</u>	<u>A216605 002</u>	Mar 08, 2023
<u>AP</u>		<u>500MG/100ML (5MG/ML)</u>	<u>A216605 003</u>	Mar 07, 2023
<u>AP</u>		<u>1GM/200ML (5MG/ML)</u>	<u>A216605 004</u>	Mar 07, 2023
<u>AP</u>	CAPLIN	<u>40MG/20ML (2MG/ML)</u>	<u>A212808 001</u>	Apr 09, 2020
<u>AP</u>		<u>100MG/20ML (5MG/ML)</u>	<u>A212808 002</u>	Apr 09, 2020
<u>AP</u>		<u>150MG/30ML (5MG/ML)</u>	<u>A212808 003</u>	Apr 09, 2020
<u>AP</u>		<u>200MG/20ML (10MG/ML)</u>	<u>A212808 004</u>	Apr 09, 2020
<u>AP</u>		<u>200MG/100ML (2MG/ML)</u>	<u>A212808 005</u>	May 23, 2024
<u>AP</u>		<u>400MG/200ML (2MG/ML)</u>	<u>A212808 006</u>	May 23, 2024
<u>AP</u>	EUGIA PHARMA	<u>40MG/20ML (2MG/ML)</u>	<u>A205612 001</u>	Jul 13, 2016
<u>AP</u>		<u>100MG/20ML (5MG/ML)</u>	<u>A205612 003</u>	Jul 13, 2016
<u>AP</u>		<u>100MG/10ML (10MG/ML)</u>	<u>A205612 006</u>	Jul 13, 2016
<u>AP</u>		<u>150MG/30ML (5MG/ML)</u>	<u>A205612 004</u>	Jul 13, 2016
<u>AP</u>		<u>150MG/20ML (7.5MG/ML)</u>	<u>A205612 005</u>	Jul 13, 2016
<u>AP</u>		<u>200MG/100ML (2MG/ML)</u>	<u>A205612 002</u>	Jul 13, 2016
<u>AP</u>		<u>200MG/20ML (10MG/ML)</u>	<u>A205612 007</u>	Jul 13, 2016
<u>AP</u>	GLAND PHARMA LTD	<u>200MG/100ML (2MG/ML)</u>	<u>A210102 001</u>	Aug 18, 2022
<u>AP</u>		<u>400MG/200ML (2MG/ML)</u>	<u>A210102 002</u>	Aug 18, 2022
<u>AP</u>	HIKMA	<u>200MG/100ML (2MG/ML)</u>	<u>A211907 001</u>	Aug 15, 2024
<u>AP</u>		<u>40MG/20ML (2MG/ML)</u>	<u>A214074 001</u>	Jul 20, 2020
<u>AP</u>		<u>150MG/30ML (5MG/ML)</u>	<u>A214074 002</u>	Jul 20, 2020
<u>AP</u>		<u>150MG/20ML (7.5MG/ML)</u>	<u>A214074 003</u>	Jul 20, 2020
<u>AP</u>		<u>200MG/20ML (10MG/ML)</u>	<u>A214074 004</u>	Jul 20, 2020
<u>AP</u>		<u>400MG/200ML (2MG/ML)</u>	<u>A211907 002</u>	Aug 15, 2024
<u>AP</u>		<u>500MG/100ML (5MG/ML)</u>	<u>A211907 003</u>	Aug 15, 2024
<u>AP</u>	INFORLIFE	<u>200MG/100ML (2MG/ML)</u>	<u>A206166 001</u>	Jun 11, 2018
<u>AP</u>		<u>400MG/200ML (2MG/ML)</u>	<u>A206166 002</u>	Jun 11, 2018
<u>AP</u>		<u>500MG/100ML (5MG/ML)</u>	<u>A206166 003</u>	Jun 11, 2018
<u>AP</u>		<u>1GM/200ML (5MG/ML)</u>	<u>A206166 004</u>	Jun 11, 2018
<u>AP</u>	KINDOS	<u>20MG/10ML (2MG/ML)</u>	<u>A218713 001</u>	Jul 30, 2024
<u>AP</u>		<u>40MG/20ML (2MG/ML)</u>	<u>A218713 002</u>	Jul 30, 2024
<u>AP</u>		<u>100MG/20ML (5MG/ML)</u>	<u>A218713 003</u>	Jul 30, 2024
<u>AP</u>		<u>100MG/10ML (10MG/ML)</u>	<u>A218713 004</u>	Jul 30, 2024
<u>AP</u>		<u>150MG/30ML (5MG/ML)</u>	<u>A218713 005</u>	Jul 30, 2024
<u>AP</u>		<u>150MG/20ML (7.5MG/ML)</u>	<u>A218713 006</u>	Jul 30, 2024
<u>AP</u>		<u>200MG/20ML (10MG/ML)</u>	<u>A218713 007</u>	Jul 30, 2024
<u>AP</u>	MYLAN LABS LTD	<u>200MG/100ML (2MG/ML)</u>	<u>A206091 001</u>	Oct 26, 2023
<u>AP</u>		<u>400MG/200ML (2MG/ML)</u>	<u>A206091 002</u>	Oct 26, 2023
<u>AP</u>	NAVINTA LLC	<u>150MG/30ML (5MG/ML)</u>	<u>A078601 002</u>	Jul 17, 2014
<u>AP</u>		<u>200MG/20ML (10MG/ML)</u>	<u>A078601 003</u>	Jul 17, 2014
<u>AP</u>	RISING	<u>200MG/100ML (2MG/ML)</u>	<u>A204636 001</u>	Mar 16, 2018
<u>AP</u>		<u>400MG/200ML (2MG/ML)</u>	<u>A204636 002</u>	Mar 16, 2018
<u>AP</u>		<u>150MG/30ML (5MG/ML)</u>	<u>A203955 001</u>	Apr 11, 2016
<u>AP</u>	SOMERSET THERAPS LLC	<u>20MG/10ML (2MG/ML)</u>	<u>A207636 001</u>	Jun 15, 2018
<u>AP</u>		<u>40MG/20ML (2MG/ML)</u>	<u>A207636 002</u>	Jun 15, 2018
<u>AP</u>		<u>100MG/20ML (5MG/ML)</u>	<u>A207636 003</u>	Jun 15, 2018
<u>AP</u>		<u>100MG/10ML (10MG/ML)</u>	<u>A207636 006</u>	Jun 15, 2018
<u>AP</u>		<u>150MG/30ML (5MG/ML)</u>	<u>A207636 004</u>	Jun 15, 2018
<u>AP</u>		<u>150MG/20ML (7.5MG/ML)</u>	<u>A207636 005</u>	Jun 15, 2018
<u>AP</u>		<u>200MG/20ML (10MG/ML)</u>	<u>A207636 007</u>	Jun 15, 2018

ROSUVASTATIN CALCIUM

TABLET; ORAL

CRESTOR

<u>AB</u>	+	<u>IPR</u>	<u>EQ 5MG BASE</u>	<u>N021366 002</u>	Aug 12, 2003
<u>AB</u>	+		<u>EQ 10MG BASE</u>	<u>N021366 003</u>	Aug 12, 2003
<u>AB</u>	+		<u>EQ 20MG BASE</u>	<u>N021366 004</u>	Aug 12, 2003
<u>AB</u>	+		<u>EQ 40MG BASE</u>	<u>N021366 005</u>	Aug 12, 2003

ROSUVASTATIN CALCIUM

<u>AB</u>	ACCORD HLTHCARE	<u>EQ 5MG BASE</u>	<u>A206434 001</u>	Oct 31, 2016
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A206434 002</u>	Oct 31, 2016
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A206434 003</u>	Oct 31, 2016

PRESCRIPTION DRUG PRODUCT LIST

ROSUVASTATIN CALCIUM

TABLET; ORAL

ROSUVASTATIN CALCIUM

<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A206434 004</u>	Oct 31, 2016
<u>AB</u>	ALKEM LABS LTD	<u>EQ 5MG BASE</u>	<u>A206465 001</u>	Mar 21, 2017
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A206465 002</u>	Mar 21, 2017
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A206465 003</u>	Mar 21, 2017
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A206465 004</u>	Mar 21, 2017
<u>AB</u>	AUROBINDO PHARMA LTD	<u>EQ 5MG BASE</u>	<u>A079170 001</u>	Jul 19, 2016
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A079170 002</u>	Jul 19, 2016
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A079170 003</u>	Jul 19, 2016
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A079170 004</u>	Jul 19, 2016
<u>AB</u>	BIOCON PHARMA	<u>EQ 5MG BASE</u>	<u>A207752 001</u>	Oct 31, 2016
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A207752 002</u>	Oct 31, 2016
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A207752 003</u>	Oct 31, 2016
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A207752 004</u>	Oct 31, 2016
<u>AB</u>	CADILA PHARMS LTD	<u>EQ 5MG BASE</u>	<u>A207453 001</u>	Nov 23, 2016
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A207453 002</u>	Nov 23, 2016
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A207453 003</u>	Nov 23, 2016
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A207453 004</u>	Nov 23, 2016
<u>AB</u>	CHANGZHOU PHARM	<u>EQ 5MG BASE</u>	<u>A207408 001</u>	Oct 31, 2016
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A207408 002</u>	Oct 31, 2016
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A207408 003</u>	Oct 31, 2016
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A207408 004</u>	Oct 31, 2016
<u>AB</u>	CHARTWELL RX	<u>EQ 5MG BASE</u>	<u>A079168 001</u>	Jul 19, 2016
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A079168 002</u>	Jul 19, 2016
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A079168 003</u>	Jul 19, 2016
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A079168 004</u>	Jul 19, 2016
<u>AB</u>	GLENMARK SPECLT	<u>EQ 5MG BASE</u>	<u>A079172 001</u>	Jul 19, 2016
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A079172 002</u>	Jul 19, 2016
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A079172 003</u>	Jul 19, 2016
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A079172 004</u>	Jul 19, 2016
<u>AB</u>	HETERO LABS LTD V	<u>EQ 5MG BASE</u>	<u>A207616 001</u>	Oct 31, 2016
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A207616 002</u>	Oct 31, 2016
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A207616 003</u>	Oct 31, 2016
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A207616 004</u>	Oct 31, 2016
<u>AB</u>	LUPIN	<u>EQ 5MG BASE</u>	<u>A205587 001</u>	Jul 31, 2017
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A205587 002</u>	Jul 31, 2017
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A205587 003</u>	Jul 31, 2017
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A205587 004</u>	Jul 31, 2017
<u>AB</u>	MACLEODS PHARMS LTD	<u>EQ 5MG BASE</u>	<u>A207296 001</u>	Nov 06, 2024
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A207296 002</u>	Nov 06, 2024
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A207296 003</u>	Nov 06, 2024
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A207296 004</u>	Nov 06, 2024
<u>AB</u>	MSN	<u>EQ 5MG BASE</u>	<u>A208898 001</u>	Nov 22, 2017
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A208898 002</u>	Nov 22, 2017
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A208898 003</u>	Nov 22, 2017
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A208898 004</u>	Nov 22, 2017
<u>AB</u>	RENATA	<u>EQ 5MG BASE</u>	<u>A207062 001</u>	Oct 31, 2016
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A207062 002</u>	Oct 31, 2016
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A207062 003</u>	Oct 31, 2016
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A207062 004</u>	Oct 31, 2016
<u>AB</u>	SCIEGEN PHARMS INC	<u>EQ 5MG BASE</u>	<u>A206381 001</u>	Apr 24, 2019
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A206381 002</u>	Apr 24, 2019
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A206381 003</u>	Apr 24, 2019
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A206381 004</u>	Apr 24, 2019
<u>AB</u>	SHANDONG	<u>EQ 5MG BASE</u>	<u>A207375 001</u>	May 07, 2019
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A207375 002</u>	May 07, 2019
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A207375 003</u>	May 07, 2019
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A207375 004</u>	May 07, 2019
<u>AB</u>	TORRENT	<u>EQ 5MG BASE</u>	<u>A201619 001</u>	Oct 31, 2016
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A201619 002</u>	Oct 31, 2016
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A201619 003</u>	Oct 31, 2016
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A201619 004</u>	Oct 31, 2016
<u>AB</u>	UMEDICA	<u>EQ 5MG BASE</u>	<u>A207626 001</u>	Apr 09, 2019
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A207626 002</u>	Apr 09, 2019
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A207626 003</u>	Apr 09, 2019
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A207626 004</u>	Apr 09, 2019
<u>AB</u>	WATSON LABS INC	<u>EQ 5MG BASE</u>	<u>A079167 001</u>	Apr 29, 2016
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A079167 002</u>	Apr 29, 2016
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A079167 003</u>	Apr 29, 2016
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A079167 004</u>	Apr 29, 2016

PRESCRIPTION DRUG PRODUCT LIST

ROSUVASTATIN CALCIUM

TABLET; ORAL

ROSUVASTATIN CALCIUM

<u>AB</u>	ZHEJIANG YONGTAI	<u>EQ 5MG BASE</u>	<u>A212059 001</u>	Nov 04, 2019
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A212059 002</u>	Nov 04, 2019
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A212059 003</u>	Nov 04, 2019
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A212059 004</u>	Nov 04, 2019

ROTIGOTINE

FILM, EXTENDED RELEASE; TRANSDERMAL

NEUPRO

+	UCB INC	1MG/24HR	N021829 004	Apr 02, 2012
+	!	2MG/24HR	N021829 001	May 09, 2007
+		3MG/24HR	N021829 005	Apr 02, 2012
+		4MG/24HR	N021829 002	May 09, 2007
+		6MG/24HR	N021829 003	May 09, 2007
+		8MG/24HR	N021829 006	Apr 02, 2012

RUBIDIUM CHLORIDE RB-82

INJECTABLE; INJECTION

CARDIOGEN-82

BRACCO

N/A

N019414 001 Dec 29, 1989

SOLUTION; INTRAVENOUS

RUBY-FILL

JUBILANT

N/A

N202153 001 Sep 30, 2016

RUCAPARIB CAMSYLATE

TABLET; ORAL

RUBRACA

+	PHARMAAND	EQ 200MG BASE	N209115 001	Dec 19, 2016
+		EQ 250MG BASE	N209115 003	May 01, 2017
+	!	EQ 300MG BASE	N209115 002	Dec 19, 2016

RUFINAMIDE

SUSPENSION; ORAL

BANZEL

<u>AB</u>	+	EISAI INC	<u>40MG/ML</u>	<u>N201367 001</u>	Mar 03, 2011
-----------	---	-----------	----------------	--------------------	--------------

RUFINAMIDE

<u>AB</u>		ALKEM LABS LTD	<u>40MG/ML</u>	<u>A213410 001</u>	Feb 23, 2021
<u>AB</u>		AUROBINDO PHARMA	<u>40MG/ML</u>	<u>A216549 001</u>	Oct 26, 2022
<u>AB</u>		BIONPHARMA	<u>40MG/ML</u>	<u>A211388 001</u>	Apr 23, 2019
<u>AB</u>		CHARTWELL RX	<u>40MG/ML</u>	<u>A214009 001</u>	Nov 07, 2022
<u>AB</u>		HETERO LABS LTD III	<u>40MG/ML</u>	<u>A216841 001</u>	Dec 05, 2023
<u>AB</u>		HIKMA	<u>40MG/ML</u>	<u>A207363 001</u>	Apr 23, 2019
<u>AB</u>		LUPIN LTD	<u>40MG/ML</u>	<u>A213457 001</u>	Dec 18, 2020

TABLET; ORAL

BANZEL

<u>AB</u>	+	EISAI INC	<u>200MG</u>	<u>N021911 002</u>	Nov 14, 2008
<u>AB</u>	+	!	<u>400MG</u>	<u>N021911 003</u>	Nov 14, 2008

RUFINAMIDE

<u>AB</u>		AUROBINDO PHARMA	<u>200MG</u>	<u>A217230 001</u>	Jun 16, 2023
<u>AB</u>			<u>400MG</u>	<u>A217230 002</u>	Jun 16, 2023
<u>AB</u>		GLENMARK PHARMS LTD	<u>200MG</u>	<u>A205075 001</u>	May 16, 2016
<u>AB</u>			<u>400MG</u>	<u>A205075 002</u>	May 16, 2016
<u>AB</u>		HETERO LABS LTD III	<u>200MG</u>	<u>A204993 001</u>	May 11, 2021
<u>AB</u>			<u>400MG</u>	<u>A204993 002</u>	May 11, 2021
<u>AB</u>		HIKMA	<u>200MG</u>	<u>A204988 001</u>	May 16, 2016
<u>AB</u>			<u>400MG</u>	<u>A204988 002</u>	May 16, 2016
<u>AB</u>		LUPIN LTD	<u>200MG</u>	<u>A204964 002</u>	Aug 17, 2022
<u>AB</u>			<u>400MG</u>	<u>A204964 003</u>	Aug 17, 2022
<u>AB</u>		MICRO LABS	<u>200MG</u>	<u>A216688 001</u>	May 15, 2023
<u>AB</u>			<u>400MG</u>	<u>A216688 002</u>	May 15, 2023
<u>AB</u>		MYLAN	<u>200MG</u>	<u>A205095 001</u>	May 16, 2016
<u>AB</u>			<u>400MG</u>	<u>A205095 002</u>	May 16, 2016
		LUPIN LTD	100MG	A204964 001	Aug 17, 2022

RUXOLITINIB PHOSPHATE

CREAM; TOPICAL

OPZELURA

+	!	INCYTE CORP	EQ 1.5% BASE	N215309 001	Sep 21, 2021
---	---	-------------	--------------	-------------	--------------

TABLET; ORAL

JAKAFI

+		INCYTE CORP	EQ 5MG BASE	N202192 001	Nov 16, 2011
+			EQ 10MG BASE	N202192 002	Nov 16, 2011
+			EQ 15MG BASE	N202192 003	Nov 16, 2011

PRESCRIPTION DRUG PRODUCT LIST

RUXOLITINIB PHOSPHATE

TABLET; ORAL

JAKAFI

+ EQ 20MG BASE
+! EQ 25MG BASEN202192 004 Nov 16, 2011
N202192 005 Nov 16, 2011SACUBITRIL; VALSARTAN

CAPSULE, PELLETS; ORAL

ENTRESTO SPRINKLE

+ NOVARTIS 6MG; 6MG
+! 15MG; 16MGN218591 001 Apr 12, 2024
N218591 002 Apr 12, 2024

TABLET; ORAL

ENTRESTO**AB** + NOVARTIS PHARMS **24MG; 26MG** **N207620 001** Jul 07, 2015
CORP**AB** + **49MG; 51MG** **N207620 002** Jul 07, 2015**AB** +! **97MG; 103MG** **N207620 003** Jul 07, 2015SACUBITRIL AND VALSARTAN**AB** ALEMBIC **24MG; 26MG** **A213682 001** May 28, 2024**AB** **49MG; 51MG** **A213682 002** May 28, 2024**AB** **97MG; 103MG** **A213682 003** May 28, 2024**AB** ALKEM LABS LTD **24MG; 26MG** **A213764 001** Sep 16, 2024**AB** **49MG; 51MG** **A213764 002** Sep 16, 2024**AB** **97MG; 103MG** **A213764 003** Sep 16, 2024**AB** BIOCON PHARMA **24MG; 26MG** **A213680 001** Aug 30, 2024**AB** **49MG; 51MG** **A213680 002** Aug 30, 2024**AB** **97MG; 103MG** **A213680 003** Aug 30, 2024**AB** CRYSTAL **24MG; 26MG** **A213605 001** May 28, 2024**AB** **49MG; 51MG** **A213605 002** May 28, 2024**AB** **97MG; 103MG** **A213605 003** May 28, 2024**AB** LAURUS **24MG; 26MG** **A213676 001** May 28, 2024**AB** **49MG; 51MG** **A213676 002** May 28, 2024**AB** **97MG; 103MG** **A213676 003** May 28, 2024**AB** MACLEODS PHARMS LTD **24MG; 26MG** **A213728 001** Oct 16, 2024**AB** **49MG; 51MG** **A213728 002** Oct 16, 2024**AB** **97MG; 103MG** **A213728 003** Oct 16, 2024**AB** MSN **24MG; 26MG** **A213748 001** Jul 24, 2024**AB** **49MG; 51MG** **A213748 002** Jul 24, 2024**AB** **97MG; 103MG** **A213748 003** Jul 24, 2024**AB** TORRENT **24MG; 26MG** **A213604 001** Aug 22, 2024**AB** **49MG; 51MG** **A213604 002** Aug 22, 2024**AB** **97MG; 103MG** **A213604 003** Aug 22, 2024**AB** ZYDUS PHARMS **24MG; 26MG** **A213719 001** Jul 09, 2024**AB** **49MG; 51MG** **A213719 002** Jul 09, 2024**AB** **97MG; 103MG** **A213719 003** Jul 09, 2024SAFINAMIDE MESYLATE

TABLET; ORAL

SAFINAMIDE MESYLATE**AB** MSN **EQ 50MG BASE** **A215978 001** Dec 31, 2024**AB** **EQ 100MG BASE** **A215978 002** Dec 31, 2024**AB** PRINSTON INC **EQ 50MG BASE** **A215739 001** Apr 25, 2024**AB** **EQ 100MG BASE** **A215739 002** Apr 25, 2024XADAGO**AB** + MDD US **EQ 50MG BASE** **N207145 001** Mar 21, 2017**AB** +! **EQ 100MG BASE** **N207145 002** Mar 21, 2017SALMETEROL XINAFOATE

POWDER; INHALATION

SEREVENT

+! GLAXOSMITHKLINE EQ 0.05MG BASE/INH

N020692 001 Sep 19, 1997

SAPROPTERIN DIHYDROCHLORIDE

POWDER; ORAL

KUVAN**AB** +! BIOMARIN PHARM **100MG/PACKET** **N205065 001** Dec 19, 2013**AB** + **500MG/PACKET** **N205065 002** Oct 27, 2015SAPROPTERIN DIHYDROCHLORIDE**AB** ANNORA PHARMA **100MG/PACKET** **A215420 001** Aug 18, 2022**AB** **500MG/PACKET** **A215420 002** Aug 18, 2022**AB** DR REDDYS **100MG/PACKET** **A209452 001** Mar 30, 2021**AB** **500MG/PACKET** **A215798 001** May 13, 2022**AB** ENDO OPERATIONS **100MG/PACKET** **A207207 001** Aug 20, 2019**AB** **500MG/PACKET** **A210027 001** Aug 20, 2019

PRESCRIPTION DRUG PRODUCT LIST

SAPROPTERIN DIHYDROCHLORIDE

TABLET; ORAL

KUVAN

AB	+ !	BIOMARIN PHARM	100MG	N022181	001	Dec 13, 2007
-----------	------------	----------------	--------------	----------------	------------	--------------

SAPROPTERIN DIHYDROCHLORIDE

AB		ANNORA PHARMA	100MG	A215534	001	Aug 23, 2022
AB		DR REDDYS	100MG	A207685	001	Jun 15, 2021
AB		ENDO OPERATIONS	100MG	A207200	001	May 10, 2019

SARECYCLINE HYDROCHLORIDE

TABLET; ORAL

SEYSARA

	+	ALMIRALL	EQ 60MG BASE	N209521	001	Oct 01, 2018
	+		EQ 100MG BASE	N209521	002	Oct 01, 2018
	+!		EQ 150MG BASE	N209521	003	Oct 01, 2018

SAXAGLIPTIN HYDROCHLORIDE

TABLET; ORAL

SAXAGLIPTIN

AB		AUROBINDO PHARMA	EQ 2.5MG BASE	A205972	001	Jul 31, 2023
AB			EQ 5MG BASE	A205972	002	Jul 31, 2023
AB		GLENMARK PHARMS LTD	EQ 2.5MG BASE	A205994	001	Jul 31, 2023
AB			EQ 5MG BASE	A205994	002	Jul 31, 2023
AB		MYLAN	EQ 2.5MG BASE	A205980	001	Jul 31, 2023
AB	!		EQ 5MG BASE	A205980	002	Jul 31, 2023
AB		SUN PHARM	EQ 2.5MG BASE	A206078	001	Jul 31, 2023
AB			EQ 5MG BASE	A206078	002	Jul 31, 2023

SCOPOLAMINE

SYSTEM; TRANSDERMAL

SCOPOLAMINE

AB		ACTAVIS LABS UT INC	1MG/72HR	A208769	001	Jan 10, 2022
AB		MYLAN TECHNOLOGIES	1MG/72HR	A203753	001	Jun 19, 2019
AB		PADAGIS US	1MG/72HR	A078830	001	Jan 30, 2015
AB		RHODES PHARMS	1MG/72HR	A215329	001	May 06, 2024
AB		RICONPHARMA LLC	1MG/72HR	A212342	001	Nov 24, 2020
AB		ZYDUS PHARMS	1MG/72HR	A217893	001	Aug 29, 2024

TRANSDERM SCOP

AB	+ !	BAXTER HLTHCARE CORP	1MG/72HR	N017874	001	
-----------	------------	----------------------	-----------------	----------------	------------	--

SECNIDAZOLE

GRANULE; ORAL

SOLOSEC

	+!	EVOFEM INC	2GM/PACKET	N209363	001	Sep 15, 2017
--	----	------------	------------	---------	-----	--------------

SECRETIN SYNTHETIC HUMAN

FOR SOLUTION; INTRAVENOUS

CHIRHOSTIM

	+!	CHIRHOCLIN	16MCG/VIAL	N021256	001	Apr 09, 2004
	+		40MCG/VIAL	N021256	002	Jun 21, 2007

SELADELPAR LYSINE

CAPSULE; ORAL

LIVDELZI

	+!	GILEAD SCIENCES INC	EQ 10MG BASE	N217899	001	Aug 14, 2024
--	----	---------------------	--------------	---------	-----	--------------

SELEGILINE

FILM, EXTENDED RELEASE; TRANSDERMAL

EMSAM

	+!	SOMERSET	6MG/24HR	N021336	001	Feb 27, 2006
	+		9MG/24HR	N021336	002	Feb 27, 2006
	+		12MG/24HR	N021336	003	Feb 27, 2006

SELEGILINE HYDROCHLORIDE

CAPSULE; ORAL

SELEGILINE HYDROCHLORIDE

AB	!	APOTEX	5MG	A075321	001	Dec 04, 1998
AB		NOVITIUM PHARMA	5MG	A075352	001	Nov 30, 1998
AB		RISING	5MG	A206803	001	Apr 02, 2019

TABLET; ORAL

SELEGILINE HYDROCHLORIDE

AB	!	APOTEX INC	5MG	A074871	001	Jun 06, 1997
AB		I3 PHARMS	5MG	A074672	001	Apr 01, 1997

PRESCRIPTION DRUG PRODUCT LIST

SELEGILINE HYDROCHLORIDE

TABLET, ORALLY DISINTEGRATING;ORAL

ZELAPAR

+! BAUSCH 1.25MG N021479 001 Jun 14, 2006

SELENIOS ACID

SOLUTION; INTRAVENOUS

SELENIOS ACID

+! AM REGENT EQ 12MCG SELENIUM/2ML (EQ 6MCG SELENIUM/ML) N209379 003 Aug 30, 2021

+! EQ 60MCG SELENIUM/ML (EQ 60MCG SELENIUM/ML) N209379 002 Jan 25, 2021

+! EQ 600MCG SELENIUM/10ML (EQ 60MCG SELENIUM/ML) N209379 001 Apr 30, 2019

SELENIUM SULFIDE

LOTION/SHAMPOO; TOPICAL

SELENIUM SULFIDE

! PADAGIS US 2.5% A089996 001 Jan 10, 1991

SELEXIPAG

POWDER; INTRAVENOUS

UPTRAVI

+! ACTELION 1.8MG/VIAL N214275 001 Jul 29, 2021

TABLET; ORAL

SELEXIPAG

AB	ALEMBIC	0.2MG	A214414 001	Oct 11, 2023
AB		0.4MG	A214414 002	Oct 11, 2023
AB		0.6MG	A214414 003	Oct 11, 2023
AB		0.8MG	A214414 004	Oct 11, 2023
AB		1.4MG	A214414 005	Oct 11, 2023
AB		1.6MG	A214414 006	Oct 11, 2023

UPTRAVI

AB	+	ACTELION	0.2MG	N207947 001	Dec 21, 2015
AB	+		0.4MG	N207947 002	Dec 21, 2015
AB	+		0.6MG	N207947 003	Dec 21, 2015
AB	+		0.8MG	N207947 004	Dec 21, 2015
AB	+		1.4MG	N207947 007	Dec 21, 2015
AB	+		1.6MG	N207947 008	Dec 21, 2015
			1MG	N207947 005	Dec 21, 2015
			1.2MG	N207947 006	Dec 21, 2015

SELINEXOR

TABLET; ORAL

XPOVIO

+ KARYOPHARM THERAPS 20MG N212306 001 Jul 03, 2019

+ 40MG N212306 002 Apr 15, 2021

+ 50MG N212306 003 Apr 15, 2021

+! 60MG N212306 004 Apr 15, 2021

SELPERCATINIB

CAPSULE; ORAL

RETEVMO

+ LOXO ONCOL ELI 40MG N213246 001 May 08, 2020

LILLY

+! 80MG N213246 002 May 08, 2020

TABLET; ORAL

RETEVMO

+ LOXO ONCOL ELI 40MG N218160 001 Apr 10, 2024

LILLY

+ 80MG N218160 002 Apr 10, 2024

+ 120MG N218160 003 Apr 10, 2024

+! 160MG N218160 004 Apr 10, 2024

SELUMETINIB SULFATE

CAPSULE; ORAL

KOSELUGO

+ ASTRAZENECA EQ 10MG BASE N213756 001 Apr 10, 2020

+! EQ 25MG BASE N213756 002 Apr 10, 2020

SEMAGLUTIDE

SOLUTION; SUBCUTANEOUS

OZEMPIC

+! NOVO 2MG/1.5ML (1.34MG/ML) N209637 001 Dec 05, 2017

+! 2MG/3ML (0.68MG/ML) N209637 004 Oct 06, 2022

+! 4MG/3ML (1.34MG/ML) N209637 002 Apr 09, 2019

+! 8MG/3ML (2.68MG/ML) N209637 003 Mar 28, 2022

PRESCRIPTION DRUG PRODUCT LIST

SEMAGLUTIDE

SOLUTION;SUBCUTANEOUS

WEGOVY

+!	NOVO	0.25MG/0.5ML (0.25MG/0.5ML)	N215256	001	Jun 04, 2021
+!		0.5MG/0.5ML (0.5MG/0.5ML)	N215256	002	Jun 04, 2021
+!		1MG/0.5ML (1MG/0.5ML)	N215256	003	Jun 04, 2021
+!		1.7MG/0.75ML (1.7MG/0.75ML)	N215256	004	Jun 04, 2021
+!		2.4MG/0.75ML (2.4MG/0.75ML)	N215256	005	Jun 04, 2021

TABLET;ORAL

RYBELSUS

+	NOVO	1.5MG	N213051	004	Dec 09, 2024
+!		3MG	N213051	001	Sep 20, 2019
+		4MG	N213051	005	Dec 09, 2024
+!		7MG	N213051	002	Sep 20, 2019
+!		9MG	N213051	006	Dec 09, 2024
+!		14MG	N213051	003	Sep 20, 2019

SERTACONAZOLE NITRATE

CREAM;TOPICAL

ERTACZO

+!	LACER PHARMA	2%	N021385	001	Dec 10, 2003
----	--------------	----	---------	-----	--------------

SERTRALINE HYDROCHLORIDE

CAPSULE;ORAL

SERTRALINE HYDROCHLORIDE

+!	ALMATICA	EQ 150MG BASE	N215133	001	Oct 04, 2021
+		EQ 200MG BASE	N215133	002	Oct 04, 2021

CONCENTRATE;ORAL

SERTRALINE HYDROCHLORIDE

AA	AUROBINDO PHARMA	EQ 20MG BASE/ML	A078861	001	Oct 31, 2008
AA	STRIDES PHARMA	EQ 20MG BASE/ML	A076934	001	Jun 30, 2006
	ZOLOFT				
AA	+! VIATRIS	EQ 20MG BASE/ML	N020990	001	Dec 07, 1999

TABLET;ORAL

SERTRALINE HYDROCHLORIDE

AB	ACCORD HLTHCARE	EQ 25MG BASE	A202825	001	Nov 07, 2014
AB		EQ 50MG BASE	A202825	002	Nov 07, 2014
AB		EQ 100MG BASE	A202825	003	Nov 07, 2014
AB	ASCENT PHARMS INC	EQ 25MG BASE	A214790	001	May 03, 2021
AB		EQ 50MG BASE	A214790	002	May 03, 2021
AB		EQ 100MG BASE	A214790	003	May 03, 2021
AB	AUROBINDO PHARMA	EQ 25MG BASE	A077206	001	Feb 06, 2007
AB		EQ 50MG BASE	A077206	002	Feb 06, 2007
AB		EQ 100MG BASE	A077206	003	Feb 06, 2007
AB	GRANULES	EQ 25MG BASE	A078403	001	Jan 08, 2008
AB		EQ 50MG BASE	A078403	002	Jan 08, 2008
AB		EQ 100MG BASE	A078403	003	Jan 08, 2008
AB	HERITAGE PHARMA AVET	EQ 25MG BASE	A076465	001	Aug 11, 2006
AB		EQ 50MG BASE	A076465	002	Aug 11, 2006
AB		EQ 100MG BASE	A076465	003	Aug 11, 2006
AB	INVAGEN PHARMS	EQ 25MG BASE	A077397	001	Feb 06, 2007
AB		EQ 50MG BASE	A077397	002	Feb 06, 2007
AB		EQ 100MG BASE	A077397	003	Feb 06, 2007
AB	LUPIN	EQ 25MG BASE	A077670	001	Feb 06, 2007
AB		EQ 50MG BASE	A077670	002	Feb 06, 2007
AB		EQ 100MG BASE	A077670	003	Feb 06, 2007
AB	OXFORD PHARMS	EQ 25MG BASE	A078175	001	Jul 21, 2010
AB		EQ 50MG BASE	A078175	002	Jul 21, 2010
AB		EQ 100MG BASE	A078175	003	Jul 21, 2010
AB	REYOUNG	EQ 25MG BASE	A078677	001	Mar 04, 2009
AB		EQ 50MG BASE	A078677	002	Mar 04, 2009
AB		EQ 100MG BASE	A078677	003	Mar 04, 2009
AB	SCIEGEN PHARMS INC	EQ 25MG BASE	A076442	001	Apr 30, 2007
AB		EQ 50MG BASE	A076442	002	Apr 30, 2007
AB		EQ 100MG BASE	A076442	003	Apr 30, 2007
AB	STRIDES PHARMA	EQ 25MG BASE	A076881	001	Feb 06, 2007
AB		EQ 50MG BASE	A076881	002	Feb 06, 2007
AB		EQ 100MG BASE	A076881	003	Feb 06, 2007
AB	VIWIT PHARM	EQ 25MG BASE	A076882	001	Feb 06, 2007
AB		EQ 50MG BASE	A076882	002	Feb 06, 2007
AB		EQ 100MG BASE	A076882	003	Feb 06, 2007

PRESCRIPTION DRUG PRODUCT LIST

SERTRALINE HYDROCHLORIDE

TABLET; ORAL

ZOLOFT

<u>AB</u>	+	VIATRIS	<u>EQ 25MG BASE</u>	<u>N019839 005</u>	Mar 06, 1996
<u>AB</u>	+		<u>EQ 50MG BASE</u>	<u>N019839 001</u>	Dec 30, 1991
<u>AB</u>	+		<u>EQ 100MG BASE</u>	<u>N019839 002</u>	Dec 30, 1991

SETMELANOTIDE ACETATE

SOLUTION; SUBCUTANEOUS

IMCIVREE

+! RHYTHM

EQ 10MG BASE/ML (EQ 10MG BASE/ML)

N213793 001 Nov 25, 2020

SEVELAMER CARBONATE

FOR SUSPENSION; ORAL

RENEVELA

<u>AB</u>	+	GENZYME	<u>800MG/PACKET</u>	<u>N022318 001</u>	Aug 12, 2009
<u>AB</u>	+		<u>2.4GM/PACKET</u>	<u>N022318 002</u>	Feb 18, 2009

SEVELAMER CARBONATE

<u>AB</u>		AUROBINDO PHARMA	<u>800MG/PACKET</u>	<u>A207624 001</u>	Jun 13, 2017
<u>AB</u>			<u>2.4GM/PACKET</u>	<u>A207624 002</u>	Jun 13, 2017
<u>AB</u>		BIONPHARMA	<u>800MG/PACKET</u>	<u>A209374 001</u>	May 17, 2021
<u>AB</u>		CHARTWELL RX	<u>800MG/PACKET</u>	<u>A217319 001</u>	Mar 02, 2023
<u>AB</u>			<u>2.4GM/PACKET</u>	<u>A217319 002</u>	Mar 02, 2023
<u>AB</u>		DR REDDYS	<u>800MG/PACKET</u>	<u>A210464 001</u>	Oct 25, 2018
<u>AB</u>			<u>2.4GM/PACKET</u>	<u>A210464 002</u>	Oct 25, 2018
<u>AB</u>		IMPAX	<u>800MG/PACKET</u>	<u>A211316 001</u>	Nov 20, 2020
<u>AB</u>			<u>2.4GM/PACKET</u>	<u>A211316 002</u>	Nov 20, 2020
<u>AB</u>		INVAGEN PHARMS	<u>2.4GM/PACKET</u>	<u>A206234 002</u>	Jul 12, 2024
<u>AB</u>			<u>800MG/PACKET</u>	<u>A206234 001</u>	Jul 12, 2024
<u>AB</u>		MACLEODS PHARMS LTD	<u>800MG/PACKET</u>	<u>A207291 001</u>	Apr 25, 2024
<u>AB</u>			<u>2.4GM/PACKET</u>	<u>A207291 002</u>	Apr 25, 2024
<u>AB</u>		STRIDES PHARMA	<u>800MG/PACKET</u>	<u>A211917 001</u>	Sep 08, 2023
<u>AB</u>			<u>2.4GM/PACKET</u>	<u>A211917 002</u>	Sep 08, 2023

TABLET; ORAL

RENEVELA

<u>AB</u>	+	SANOFI	<u>800MG</u>	<u>N022127 001</u>	Oct 19, 2007
-----------	---	--------	--------------	--------------------	--------------

SEVELAMER CARBONATE

<u>AB</u>		AMNEAL PHARMS CO	<u>800MG</u>	<u>A207288 001</u>	Nov 28, 2017
<u>AB</u>		ARTHUR GRP	<u>800MG</u>	<u>A200959 001</u>	Mar 20, 2018
<u>AB</u>		AUROBINDO PHARMA	<u>800MG</u>	<u>A207179 001</u>	Jul 17, 2017
<u>AB</u>		DR REDDYS	<u>800MG</u>	<u>A206094 001</u>	Sep 29, 2017
<u>AB</u>		INVAGEN PHARMS	<u>800MG</u>	<u>A203860 001</u>	Oct 26, 2017
<u>AB</u>		MACLEODS PHARMS LTD	<u>800MG</u>	<u>A206100 001</u>	Apr 19, 2023
<u>AB</u>		MICRO LABS	<u>800MG</u>	<u>A215537 001</u>	Feb 07, 2022
<u>AB</u>		RISING	<u>800MG</u>	<u>A204451 001</u>	Nov 29, 2018
<u>AB</u>		SHANDONG XINHUA	<u>800MG</u>	<u>A215998 001</u>	Oct 24, 2024
<u>AB</u>		STRIDES PHARMA	<u>800MG</u>	<u>A211915 001</u>	May 09, 2024
<u>AB</u>		ZYDUS PHARMS	<u>800MG</u>	<u>A207759 001</u>	Aug 11, 2020

SEVELAMER HYDROCHLORIDE

TABLET; ORAL

RENAGEL

<u>AB</u>	+	GENZYME	<u>400MG</u>	<u>N021179 001</u>	Jul 12, 2000
<u>AB</u>	+		<u>800MG</u>	<u>N021179 002</u>	Jul 12, 2000

SEVELAMER HYDROCHLORIDE

<u>AB</u>		AUROBINDO PHARMA LTD	<u>400MG</u>	<u>A212599 001</u>	Jul 11, 2023
<u>AB</u>			<u>800MG</u>	<u>A212599 002</u>	Jul 11, 2023
<u>AB</u>		GLENMARK PHARMS LTD	<u>400MG</u>	<u>A204724 001</u>	Feb 08, 2019
<u>AB</u>			<u>800MG</u>	<u>A204724 002</u>	Feb 08, 2019
<u>AB</u>		LUPIN LTD	<u>400MG</u>	<u>A213145 001</u>	Jun 16, 2021
<u>AB</u>			<u>800MG</u>	<u>A213145 002</u>	Jun 16, 2021
<u>AB</u>		MACLEODS PHARMS LTD	<u>400MG</u>	<u>A206883 001</u>	May 26, 2023
<u>AB</u>			<u>800MG</u>	<u>A206883 002</u>	May 26, 2023

SEVOFLURANE

LIQUID; INHALATION

SEVOFLURANE

<u>AN</u>		BAXTER HLTHCARE	<u>100%</u>	<u>A075895 001</u>	Jul 02, 2002
<u>AN</u>		HALOCARBON PRODS	<u>100%</u>	<u>A078650 001</u>	Nov 19, 2007
<u>AN</u>		SHANDONG	<u>100%</u>	<u>A214382 001</u>	Aug 18, 2023
<u>AN</u>		SHANGHAI HENGRUI	<u>100%</u>	<u>A203793 001</u>	Nov 03, 2015
		<u>SOJOURN</u>			
<u>AN</u>		PIRAMAL CRITICAL	<u>100%</u>	<u>A077867 001</u>	May 02, 2007

PRESCRIPTION DRUG PRODUCT LIST

SEVOFLURANE

LIQUID; INHALATION

ULTANE

AN	+! ABBVIE	100%	N020478	001	Jun 07, 1995
-----------	------------------	-------------	----------------	------------	--------------

SILDENAFIL CITRATE

FOR SUSPENSION; ORAL

REVATIO

AB	+! VIATRIS	EQ 10MG BASE/ML	N203109	001	Aug 30, 2012
-----------	-------------------	------------------------	----------------	------------	--------------

SILDENAFIL CITRATE

AB	AJANTA PHARMA LTD	EQ 10MG BASE/ML	A212883	001	Nov 27, 2019
AB	ALKEM LABS LTD	EQ 10MG BASE/ML	A212440	001	Nov 27, 2019
AB	AMNEAL PHARMS	EQ 10MG BASE/ML	A211092	001	Nov 27, 2019
AB	AUROBINDO PHARMA LTD	EQ 10MG BASE/ML	A214773	001	Dec 23, 2022
AB	HETERO LABS LTD V	EQ 10MG BASE/ML	A213014	001	May 11, 2021
AB	INVAGEN PHARMS	EQ 10MG BASE/ML	A213041	001	Aug 06, 2020
AB	LUPIN LTD	EQ 10MG BASE/ML	A211638	001	Mar 23, 2022
AB	MSN	EQ 10MG BASE/ML	A214641	001	Feb 08, 2022
AB	NOVITIUM PHARMA	EQ 10MG BASE/ML	A212260	001	May 31, 2019
AB	SOMERSET THERAPS LLC	EQ 10MG BASE/ML	A213814	001	Apr 29, 2021
AB	TARO	EQ 10MG BASE/ML	A215522	001	Nov 16, 2021
AB	TEVA PHARMS USA	EQ 10MG BASE/ML	A211534	001	May 29, 2020
AB	ZYDUS LIFESCIENCES	EQ 10MG BASE/ML	A215708	001	Sep 29, 2022

SOLUTION; INTRAVENOUS

REVATIO

AP	+! VIATRIS	EQ 10MG BASE/12.5ML (EQ 0.8MG BASE/ML)	N022473	001	Nov 18, 2009
-----------	-------------------	---	----------------	------------	--------------

SILDENAFIL CITRATE

AP	EUGIA PHARMA	EQ 10MG BASE/12.5ML (EQ 0.8MG BASE/ML)	A203988	001	Apr 01, 2015
-----------	--------------	---	----------------	------------	--------------

TABLET; ORAL

REVATIO

AB	+! VIATRIS	EQ 20MG BASE	N021845	001	Jun 03, 2005
-----------	-------------------	---------------------	----------------	------------	--------------

SILDENAFIL CITRATE

AB	AJANTA PHARMA LTD	EQ 20MG BASE	A210394	001	May 04, 2018
AB		EQ 25MG BASE	A206401	001	Oct 12, 2018
AB		EQ 50MG BASE	A206401	002	Oct 12, 2018
AB		EQ 100MG BASE	A206401	003	Oct 12, 2018
AB	AMNEAL PHARMS	EQ 20MG BASE	A202025	001	Feb 28, 2013
AB	AMNEAL PHARMS NY	EQ 25MG BASE	A202023	001	Jun 27, 2018
AB		EQ 50MG BASE	A202023	002	Jun 27, 2018
AB		EQ 100MG BASE	A202023	003	Jun 27, 2018
AB	APPCO	EQ 25MG BASE	A207178	001	Mar 02, 2020
AB		EQ 50MG BASE	A207178	002	Mar 02, 2020
AB		EQ 100MG BASE	A207178	003	Mar 02, 2020
AB	AUROBINDO PHARMA LTD	EQ 20MG BASE	A203963	001	Nov 18, 2015
AB		EQ 25MG BASE	A203962	001	Jun 11, 2018
AB		EQ 50MG BASE	A203962	002	Jun 11, 2018
AB		EQ 100MG BASE	A203962	003	Jun 11, 2018
AB	CADILA PHARMS LTD	EQ 25MG BASE	A218045	001	Nov 28, 2023
AB		EQ 50MG BASE	A218045	002	Nov 28, 2023
AB		EQ 100MG BASE	A218045	003	Nov 28, 2023
AB	CHARTWELL RX	EQ 20MG BASE	A202598	001	Nov 06, 2012
AB		EQ 25MG BASE	A212051	001	Mar 22, 2019
AB		EQ 50MG BASE	A212051	002	Mar 22, 2019
AB		EQ 100MG BASE	A212051	003	Mar 22, 2019
AB	HETERO LABS LTD V	EQ 20MG BASE	A203623	001	Nov 26, 2014
AB		EQ 25MG BASE	A202659	001	Jun 11, 2018
AB		EQ 50MG BASE	A202659	002	Jun 11, 2018
AB		EQ 100MG BASE	A202659	003	Jun 11, 2018
AB	JUBILANT GENERICS	EQ 25MG BASE	A201156	001	Dec 20, 2024
AB		EQ 50MG BASE	A201156	002	Dec 20, 2024
AB		EQ 100MG BASE	A201156	003	Dec 20, 2024
AB	MACLEODS PHARMS LTD	EQ 20MG BASE	A203814	001	Dec 17, 2013
AB		EQ 25MG BASE	A202255	001	Apr 19, 2023
AB		EQ 50MG BASE	A202255	002	Apr 19, 2023
AB		EQ 100MG BASE	A202255	003	Apr 19, 2023
AB	MYLAN	EQ 25MG BASE	A201171	001	Mar 25, 2019
AB		EQ 50MG BASE	A201171	002	Mar 25, 2019
AB		EQ 100MG BASE	A201171	003	Mar 25, 2019
AB	NOVITIUM PHARMA	EQ 25MG BASE	A216383	001	Aug 29, 2023
AB		EQ 50MG BASE	A216383	002	Aug 29, 2023

PRESCRIPTION DRUG PRODUCT LIST

SILDENAFIL CITRATE

TABLET; ORAL

SILDENAFIL CITRATE

<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A216383 003</u>	Aug 29, 2023
<u>AB</u>	REYOUNG	<u>EQ 25MG BASE</u>	<u>A208494 001</u>	Jun 12, 2020
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A208494 002</u>	Jun 12, 2020
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A208494 003</u>	Jun 12, 2020
<u>AB</u>	RUBICON	<u>EQ 20MG BASE</u>	<u>A204883 001</u>	Jun 20, 2016
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>A204882 001</u>	Jun 11, 2018
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A204882 002</u>	Jun 11, 2018
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A204882 003</u>	Jun 11, 2018
<u>AB</u>	SUNSHINE	<u>EQ 25MG BASE</u>	<u>A213032 001</u>	Jun 11, 2020
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A213032 002</u>	Jun 11, 2020
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A213032 003</u>	Jun 11, 2020
<u>AB</u>	TEVA	<u>EQ 25MG BASE</u>	<u>A077342 001</u>	Mar 09, 2016
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A077342 002</u>	Mar 09, 2016
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A077342 003</u>	Mar 09, 2016
<u>AB</u>	TEVA PHARMS	<u>EQ 20MG BASE</u>	<u>A078380 001</u>	Jan 07, 2013
<u>AB</u>	TORRENT	<u>EQ 25MG BASE</u>	<u>A091448 001</u>	Jun 11, 2018
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A091448 002</u>	Jun 11, 2018
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A091448 003</u>	Jun 11, 2018
<u>AB</u>	TORRENT PHARMS LTD	<u>EQ 20MG BASE</u>	<u>A091479 001</u>	Nov 06, 2012
<u>AB</u>	WATSON LABS INC	<u>EQ 20MG BASE</u>	<u>A202503 001</u>	Nov 06, 2012

VIAGRA

<u>AB</u>	+ VIATRIS	<u>EQ 25MG BASE</u>	<u>N020895 001</u>	Mar 27, 1998
<u>AB</u>	+	<u>EQ 50MG BASE</u>	<u>N020895 002</u>	Mar 27, 1998
<u>AB</u>	+	<u>EQ 100MG BASE</u>	<u>N020895 003</u>	Mar 27, 1998
	SILDENAFIL CITRATE			
BX	UMEDICA	EQ 25MG BASE	A209302 003	Jun 15, 2021
BX		EQ 50MG BASE	A209302 001	Aug 25, 2020
BX		EQ 100MG BASE	A209302 002	Aug 25, 2020

SILODOSIN

CAPSULE; ORAL

RAPAFLO

<u>AB</u>	+! ABBVIE	<u>4MG</u>	<u>N022206 001</u>	Oct 08, 2008
<u>AB</u>	+	<u>8MG</u>	<u>N022206 002</u>	Oct 08, 2008

SILODOSIN

<u>AB</u>	AJANTA PHARMA LTD	<u>4MG</u>	<u>A211060 001</u>	Dec 03, 2018
<u>AB</u>		<u>8MG</u>	<u>A211060 002</u>	Dec 03, 2018
<u>AB</u>	AMNEAL PHARMS CO	<u>4MG</u>	<u>A209745 001</u>	Dec 03, 2018
<u>AB</u>		<u>8MG</u>	<u>A209745 002</u>	Dec 03, 2018
<u>AB</u>	AUROBINDO PHARMA LTD	<u>4MG</u>	<u>A210626 001</u>	Dec 10, 2018
<u>AB</u>		<u>8MG</u>	<u>A210626 002</u>	Dec 10, 2018
<u>AB</u>	CHARTWELL RX	<u>4MG</u>	<u>A204726 001</u>	Mar 31, 2017
<u>AB</u>		<u>8MG</u>	<u>A204726 002</u>	Mar 31, 2017
<u>AB</u>	CREEKWOOD PHARMS	<u>4MG</u>	<u>A213230 001</u>	Jan 03, 2022
<u>AB</u>		<u>8MG</u>	<u>A213230 002</u>	Jan 03, 2022
<u>AB</u>	HETERO LABS LTD V	<u>4MG</u>	<u>A204793 001</u>	Feb 14, 2020
<u>AB</u>		<u>8MG</u>	<u>A204793 002</u>	Feb 14, 2020
<u>AB</u>	LUPIN LTD	<u>4MG</u>	<u>A206541 001</u>	Dec 03, 2018
<u>AB</u>		<u>8MG</u>	<u>A206541 002</u>	Dec 03, 2018
<u>AB</u>	MACLEODS PHARMS LTD	<u>4MG</u>	<u>A211166 001</u>	Dec 03, 2018
<u>AB</u>		<u>8MG</u>	<u>A211166 002</u>	Dec 03, 2018
<u>AB</u>	MSN	<u>4MG</u>	<u>A210687 001</u>	Dec 03, 2018
<u>AB</u>		<u>8MG</u>	<u>A210687 002</u>	Dec 03, 2018
<u>AB</u>	PRINSTON INC	<u>4MG</u>	<u>A209029 001</u>	Jan 04, 2022
<u>AB</u>		<u>8MG</u>	<u>A209029 002</u>	Jan 04, 2022

SILVER SULFADIAZINE

CREAM; TOPICAL

SILVADENE

<u>AB</u>	+! KING PHARMS LLC	<u>1%</u>	<u>N017381 001</u>	
	<u>SSD</u>			
<u>AB</u>	+! DR REDDYS	<u>1%</u>	<u>N018578 001</u>	Feb 25, 1982
	<u>THERMAZENE</u>			
<u>AB</u>	THEPHARMANETWORK LLC	<u>1%</u>	<u>N018810 001</u>	Dec 23, 1985

PRESCRIPTION DRUG PRODUCT LIST

SIMVASTATIN

SUSPENSION; ORAL

FLOLIPID

+ SALERNO PHARMS

20MG/5ML

N206679 001 Apr 21, 2016

+!

40MG/5ML

N206679 002 Apr 21, 2016

TABLET; ORAL

SIMVASTATIN

<u>AB</u>	ACCORD HLTHCARE	<u>5MG</u>	<u>A078155 005</u>	Apr 05, 2013
<u>AB</u>		<u>10MG</u>	<u>A078155 002</u>	Feb 26, 2008
<u>AB</u>		<u>20MG</u>	<u>A078155 003</u>	Feb 26, 2008
<u>AB</u>		<u>40MG</u>	<u>A078155 004</u>	Feb 26, 2008
<u>AB</u>		<u>80MG</u>	<u>A078155 001</u>	Feb 26, 2008
<u>AB</u>	AUROBINDO PHARMA	<u>5MG</u>	<u>A077691 001</u>	Dec 20, 2006
<u>AB</u>		<u>10MG</u>	<u>A077691 002</u>	Dec 20, 2006
<u>AB</u>		<u>20MG</u>	<u>A077691 003</u>	Dec 20, 2006
<u>AB</u>		<u>40MG</u>	<u>A077691 004</u>	Dec 20, 2006
<u>AB</u>		<u>80MG</u>	<u>A077691 005</u>	Dec 20, 2006
<u>AB</u>	BIOCON PHARMA	<u>5MG</u>	<u>A078034 001</u>	Dec 20, 2006
<u>AB</u>		<u>10MG</u>	<u>A078034 002</u>	Dec 20, 2006
<u>AB</u>		<u>20MG</u>	<u>A078034 003</u>	Dec 20, 2006
<u>AB</u>		<u>40MG</u>	<u>A078034 004</u>	Dec 20, 2006
<u>AB</u>		<u>80MG</u>	<u>A078034 005</u>	Dec 20, 2006
<u>AB</u>	DR REDDYS LABS INC	<u>5MG</u>	<u>A077752 005</u>	Jan 23, 2008
<u>AB</u>		<u>10MG</u>	<u>A077752 001</u>	Dec 20, 2006
<u>AB</u>		<u>20MG</u>	<u>A077752 002</u>	Dec 20, 2006
<u>AB</u>		<u>40MG</u>	<u>A077752 003</u>	Dec 20, 2006
<u>AB</u>		<u>80MG</u>	<u>A077752 004</u>	Dec 20, 2006
<u>AB</u>	HETERO LABS LTD III	<u>5MG</u>	<u>A200895 001</u>	Nov 25, 2014
<u>AB</u>		<u>10MG</u>	<u>A200895 002</u>	Nov 25, 2014
<u>AB</u>		<u>20MG</u>	<u>A200895 003</u>	Nov 25, 2014
<u>AB</u>		<u>40MG</u>	<u>A200895 004</u>	Nov 25, 2014
<u>AB</u>		<u>80MG</u>	<u>A200895 005</u>	Nov 25, 2014
<u>AB</u>	INVAGEN PHARMS	<u>10MG</u>	<u>A206557 001</u>	Nov 13, 2017
<u>AB</u>		<u>20MG</u>	<u>A206557 002</u>	Nov 13, 2017
<u>AB</u>		<u>40MG</u>	<u>A206557 003</u>	Nov 13, 2017
<u>AB</u>		<u>80MG</u>	<u>A206557 004</u>	Nov 13, 2017
<u>AB</u>	LUPIN	<u>5MG</u>	<u>A078103 005</u>	Apr 14, 2009
<u>AB</u>		<u>10MG</u>	<u>A078103 001</u>	May 11, 2007
<u>AB</u>		<u>20MG</u>	<u>A078103 002</u>	May 11, 2007
<u>AB</u>		<u>40MG</u>	<u>A078103 003</u>	May 11, 2007
<u>AB</u>	!	<u>80MG</u>	<u>A078103 004</u>	May 11, 2007
<u>AB</u>	MICRO LABS	<u>5MG</u>	<u>A090383 001</u>	Sep 16, 2011
<u>AB</u>		<u>10MG</u>	<u>A090383 002</u>	Sep 16, 2011
<u>AB</u>		<u>20MG</u>	<u>A090383 003</u>	Sep 16, 2011
<u>AB</u>		<u>40MG</u>	<u>A090383 004</u>	Sep 16, 2011
<u>AB</u>		<u>80MG</u>	<u>A090383 005</u>	Sep 16, 2011
<u>AB</u>	OXFORD PHARMS	<u>5MG</u>	<u>A078735 001</u>	Aug 30, 2010
<u>AB</u>		<u>10MG</u>	<u>A078735 002</u>	Aug 30, 2010
<u>AB</u>		<u>20MG</u>	<u>A078735 003</u>	Aug 30, 2010
<u>AB</u>		<u>40MG</u>	<u>A078735 004</u>	Aug 30, 2010
<u>AB</u>		<u>80MG</u>	<u>A078735 005</u>	Aug 30, 2010
<u>AB</u>	YILING	<u>10MG</u>	<u>A211394 001</u>	Nov 27, 2024
<u>AB</u>		<u>20MG</u>	<u>A211394 002</u>	Nov 27, 2024
<u>AB</u>		<u>40MG</u>	<u>A211394 003</u>	Nov 27, 2024
<u>AB</u>		<u>80MG</u>	<u>A211394 004</u>	Nov 27, 2024
<u>AB</u>	ZYDUS PHARMS USA	<u>5MG</u>	<u>A077837 001</u>	Dec 20, 2006
<u>AB</u>		<u>10MG</u>	<u>A077837 002</u>	Dec 20, 2006
<u>AB</u>		<u>20MG</u>	<u>A077837 003</u>	Dec 20, 2006
<u>AB</u>		<u>40MG</u>	<u>A077837 004</u>	Dec 20, 2006
<u>AB</u>		<u>80MG</u>	<u>A077837 005</u>	Dec 20, 2006
<u>ZOCOR</u>				
<u>AB</u>	+ ORGANON	<u>5MG</u>	<u>N019766 001</u>	Dec 23, 1991
<u>AB</u>	+	<u>10MG</u>	<u>N019766 002</u>	Dec 23, 1991
<u>AB</u>	+	<u>20MG</u>	<u>N019766 003</u>	Dec 23, 1991
<u>AB</u>	+	<u>40MG</u>	<u>N019766 004</u>	Dec 23, 1991

SINCALIDE

POWDER; INTRAVENOUS

KINEVAC

AP +! BRACCO **0.005MG/VIAL** **N017697 001**

SINCALIDE

AP +! MAIA PHARMS INC **0.005MG/VIAL** **N210850 001** Nov 22, 2022

PRESCRIPTION DRUG PRODUCT LIST

SINECATECHINS

OINTMENT; TOPICAL

VEREGEN

+! ANI PHARMS 15% N021902 001 Oct 31, 2006

SIPONIMOD

TABLET; ORAL

MAYZENT

+ NOVARTIS EQ 0.25MG BASE N209884 001 Mar 26, 2019

+ EQ 1MG BASE N209884 003 Aug 24, 2021

+! EQ 2MG BASE N209884 002 Mar 26, 2019

SIROLIMUS

GEL; TOPICAL

HYFTOR

+! NOBELPHARMA 0.2% N213478 001 Mar 22, 2022

POWDER; INTRAVENOUS

FYARRO

+! AADI 100MG/VIAL N213312 001 Nov 22, 2021

SOLUTION; ORAL

RAPAMUNE**AA** +! PF PRISM CV 1MG/ML **N021083 001** Sep 15, 1999SIROLIMUS**AA** AMNEAL 1MG/ML **A211212 001** Oct 18, 2019**AA** APOTEX 1MG/ML **A211406 001** Oct 22, 2019**AA** HETERO LABS LTD V 1MG/ML **A215080 001** Sep 16, 2024**AA** MSN 1MG/ML **A216728 001** Jan 19, 2023**AA** NOVITIUM PHARMA 1MG/ML **A211040 001** Jan 28, 2019

TABLET; ORAL

RAPAMUNE**AB** + PF PRISM CV 0.5MG **N021110 004** Jan 25, 2010**AB** + 1MG **N021110 001** Aug 25, 2000**AB** +! 2MG **N021110 002** Aug 22, 2002SIROLIMUS**AB** ALKEM LABS LTD 0.5MG **A214753 001** Mar 12, 2021**AB** 1MG **A214753 002** Mar 12, 2021**AB** 2MG **A214753 003** Mar 12, 2021**AB** DR REDDYS 1MG **A201578 001** Oct 27, 2014**AB** 2MG **A201578 002** Oct 27, 2014**AB** GLENMARK PHARMS LTD 0.5MG **A208691 001** Oct 16, 2020**AB** 1MG **A208691 002** Oct 16, 2020**AB** 2MG **A208691 003** Oct 16, 2020**AB** ZYDUS PHARMS 1MG **A201676 001** Feb 15, 2023**AB** 2MG **A201676 002** Feb 15, 2023**AB** 0.5MG **A201676 003** Jan 08, 2014SITAGLIPTIN

TABLET; ORAL

ZITUVIO

+ ZYDUS LIFESCIENCES 25MG N211566 001 Oct 18, 2023

+ 50MG N211566 002 Oct 18, 2023

+! 100MG N211566 003 Oct 18, 2023

SITAGLIPTIN PHOSPHATE

TABLET; ORAL

JANUVIA

+ MERCK SHARP DOHME EQ 25MG BASE N021995 001 Oct 16, 2006

+ EQ 50MG BASE N021995 002 Oct 16, 2006

+! EQ 100MG BASE N021995 003 Oct 16, 2006

SODIUM ACETATE

SOLUTION; INTRAVENOUS

SODIUM ACETATE**AP** ! FRESENIUS KABI USA 400MEQ/100ML (4MEQ/ML) **A206687 001** Oct 30, 2017**AP** HIKMA 400MEQ/20ML (2MEQ/ML) **A216920 001** Mar 15, 2024**AP** 100MEQ/50ML (2MEQ/ML) **A216920 002** Mar 15, 2024**AP** 200MEQ/100ML (2MEQ/ML) **A216920 003** Mar 15, 2024**AP** +! HOSPIRA 400MEQ/20ML (2MEQ/ML) **N018893 001** May 04, 1983**AP** +! 100MEQ/50ML (2MEQ/ML) **N018893 002** Sep 02, 2014**AP** +! 200MEQ/100ML (2MEQ/ML) **N018893 003** Sep 02, 2014**AP** MILLA PHARMS 400MEQ/20ML (2MEQ/ML) **A214805 004** Jan 22, 2024**AP** 100MEQ/50ML (2MEQ/ML) **A214805 001** May 04, 2021**AP** 200MEQ/100ML (2MEQ/ML) **A214805 003** May 04, 2021**AP** 400MEQ/100ML (4MEQ/ML) **A214805 002** Aug 25, 2023

PRESCRIPTION DRUG PRODUCT LIST

SODIUM BENZOATE; SODIUM PHENYLACETATE

SOLUTION; INTRAVENOUS

SODIUM PHENYLACETATE AND SODIUM BENZOATE

<u>AP</u>	!	AILEX PHARMS LLC	<u>10%;10% (5GM/50ML;5GM/50ML)</u>	<u>A207096 001</u>	Feb 24, 2016
<u>AP</u>		MAIA PHARMS INC	<u>10%;10% (5GM/50ML;5GM/50ML)</u>	<u>A208521 001</u>	May 08, 2017
<u>AP</u>	+		<u>10%;10% (2GM/20ML;2GM/20ML)</u>	<u>N215025 001</u>	Jun 10, 2021
<u>AP</u>		NAVINTA LLC	<u>10%;10% (5GM/50ML;5GM/50ML)</u>	<u>A205880 001</u>	Aug 04, 2016
<u>AP</u>			<u>10%;10% (2GM/20ML; 2GM/20ML)</u>	<u>A217526 001</u>	Jul 14, 2023

SODIUM BICARBONATE

INJECTABLE; INJECTION

SODIUM BICARBONATE

<u>AP</u>	!	EXELA PHARMA	<u>0.5MEQ/ML</u>	<u>A211091 001</u>	Jun 20, 2019
<u>AP</u>	!		<u>0.9MEQ/ML</u>	<u>A211091 002</u>	Jun 20, 2019
<u>AP</u>	!		<u>1MEQ/ML</u>	<u>A211091 003</u>	Jun 20, 2019
<u>AP</u>		HOSPIRA	<u>0.5MEQ/ML</u>	<u>A202679 001</u>	Mar 07, 2017
<u>AP</u>	!		<u>0.5MEQ/ML</u>	<u>A202981 001</u>	Mar 04, 2016
<u>AP</u>			<u>0.9MEQ/ML</u>	<u>A202494 001</u>	Mar 06, 2017
<u>AP</u>			<u>1MEQ/ML</u>	<u>A202432 001</u>	Sep 26, 2017
<u>AP</u>			<u>1MEQ/ML</u>	<u>A202494 002</u>	Mar 06, 2017
<u>AP</u>		INTL MEDICATION SYS	<u>1MEQ/ML</u>	<u>A203449 001</u>	Sep 19, 2017
<u>AP</u>		LONG GROVE PHARMS	<u>1MEQ/ML</u>	<u>A216042 001</u>	Feb 27, 2024
<u>AP</u>		OMNIVIUM PHARMS	<u>1MEQ/ML</u>	<u>A216364 001</u>	Jun 15, 2023
<u>AP</u>		STERISCIENCE	<u>1MEQ/ML</u>	<u>A217594 001</u>	Jun 28, 2023
<u>AP</u>		HOSPIRA	1MEQ/ML	A202495 001	Mar 06, 2017

SODIUM CHLORIDE

INJECTABLE; INJECTION

SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER

<u>AP</u>	+	B BRAUN	<u>450MG/100ML</u>	<u>N019635 001</u>	Mar 09, 1988
<u>AP</u>	+	BAXTER HLTHCARE	<u>450MG/100ML</u>	<u>N018016 001</u>	
<u>AP</u>		FRESENIUS KABI USA	<u>450MG/100ML</u>	<u>A208122 001</u>	Jul 23, 2018
<u>AP</u>		HOSPIRA	<u>450MG/100ML</u>	<u>N019759 001</u>	Jun 08, 1988
<u>AP</u>	+	ICU MEDICAL INC	<u>450MG/100ML</u>	<u>N018090 001</u>	

SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

<u>AP</u>	+	B BRAUN	<u>900MG/100ML</u>	<u>N017464 001</u>	
<u>AP</u>	+		<u>900MG/100ML</u>	<u>N019635 002</u>	Mar 09, 1988
<u>AP</u>		FRESENIUS KABI USA	<u>900MG/100ML</u>	<u>A207310 001</u>	Sep 19, 2017
<u>AP</u>		FRESENIUS MEDCL	<u>900MG/100ML</u>	<u>A078177 001</u>	Apr 12, 2007
<u>AP</u>		HAEMONETICS	<u>900MG/100ML</u>	<u>A076316 001</u>	Oct 27, 2004
<u>AP</u>	+	HOSPIRA	<u>900MG/100ML</u>	<u>N019480 001</u>	Sep 17, 1985
<u>AP</u>	+	ICU MEDICAL INC	<u>900MG/100ML</u>	<u>N016366 001</u>	
<u>AP</u>		LABORATORIOS GRIFOLS	<u>900MG/100ML</u>	<u>A207956 001</u>	May 25, 2017

SODIUM CHLORIDE 23.4%

<u>AP</u>		FRESENIUS KABI USA	<u>234MG/ML</u>	<u>A212248 001</u>	Apr 28, 2021
<u>AP</u>			<u>234MG/ML</u>	<u>A217796 001</u>	Jul 11, 2023

SODIUM CHLORIDE 3% IN PLASTIC CONTAINER

<u>AP</u>		B BRAUN	<u>3GM/100ML</u>	<u>N019635 003</u>	Mar 09, 1988
<u>AP</u>	+	BAXTER HLTHCARE	<u>3GM/100ML</u>	<u>N019022 001</u>	Nov 01, 1983
<u>AP</u>		FRESENIUS KABI USA	<u>3GM/100ML</u>	<u>A209476 001</u>	Mar 13, 2019

SODIUM CHLORIDE 5% IN PLASTIC CONTAINER

<u>AP</u>		B BRAUN	<u>5GM/100ML</u>	<u>N019635 004</u>	Mar 09, 1988
<u>AP</u>	+	BAXTER HLTHCARE	<u>5GM/100ML</u>	<u>N019022 002</u>	Nov 01, 1983

SODIUM CHLORIDE 0.9%

+	!	MEDEFIL INC	9MG/ML (9MG/ML)	N202832 001	Jan 06, 2012
	+		18MG/2ML (9MG/ML)	N202832 002	Jan 06, 2012
	+		22.5MG/2.5ML (9MG/ML)	N202832 003	Jan 06, 2012
	+		27MG/3ML (9MG/ML)	N202832 004	Jan 06, 2012
	+		45MG/5ML (9MG/ML)	N202832 005	Jan 06, 2012
	+		90MG/10ML (9MG/ML)	N202832 006	Jan 06, 2012

SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

+		LIEBEL-FLARSHEIM	1.125GM/125ML (9MG/ML)	N021569 002	Jul 27, 2006
---	--	------------------	------------------------	-------------	--------------

SOLUTION; INJECTION

BACTERIOSTATIC SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

<u>AP</u>		FRESENIUS KABI USA	<u>90MG/10ML (9MG/ML)</u>	<u>A088911 002</u>	May 17, 1985
<u>AP</u>			<u>270MG/30ML (9MG/ML)</u>	<u>A088911 001</u>	Feb 07, 1985
<u>AP</u>	+	HOSPIRA	<u>90MG/10ML (9MG/ML)</u>	<u>N018800 001</u>	Oct 29, 1982
<u>AP</u>	+		<u>180MG/20ML (9MG/ML)</u>	<u>N018800 002</u>	Oct 29, 1982
<u>AP</u>	+		<u>270MG/30ML (9MG/ML)</u>	<u>N018800 003</u>	Oct 29, 1982

SODIUM CHLORIDE 0.9%

<u>AP</u>		HIKMA	<u>90MG/10ML (9MG/ML)</u>	<u>A201833 002</u>	Jan 07, 2015
<u>AP</u>		NEXUS PHARMS	<u>90MG/10ML (9MG/ML)</u>	<u>A217535 001</u>	Aug 23, 2023
<u>AP</u>			<u>180MG/20ML (9MG/ML)</u>	<u>A217535 002</u>	Aug 23, 2023

PRESCRIPTION DRUG PRODUCT LIST

SODIUM CHLORIDE

SOLUTION; INJECTION

SODIUM CHLORIDE 0.9%

<u>AP</u>	SPECTRA MDCL DEVICES	<u>45MG/5ML (9MG/ML)</u>	<u>A206171 001</u>	Jul 21, 2017
<u>AP</u>		<u>90MG/10ML (9MG/ML)</u>	<u>A206171 002</u>	Jul 21, 2017
<u>SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER</u>				
<u>AP</u>	+!	BAXTER HLTHCARE	<u>450MG/50ML (9MG/ML)</u>	<u>N016677 003</u>
<u>AP</u>	+		<u>450MG/50ML (9MG/ML)</u>	<u>N020178 002</u> Dec 07, 1992
<u>AP</u>	+!		<u>900MG/100ML</u>	<u>N016677 001</u>
<u>AP</u>	+!		<u>900MG/100ML</u>	<u>N020178 001</u> Dec 07, 1992
<u>AP</u>	!	FRESENIUS KABI USA	<u>90MG/10ML (9MG/ML)</u>	<u>A088912 001</u> Jan 10, 1985
<u>AP</u>	!		<u>180MG/20ML (9MG/ML)</u>	<u>A088912 002</u> Jan 10, 1985
<u>AP</u>	!		<u>270MG/30ML (9MG/ML)</u>	<u>A088912 003</u> Jan 10, 1985
<u>AP</u>	+!	HOSPIRA	<u>90MG/10ML (9MG/ML)</u>	<u>N018803 001</u> Oct 29, 1982
<u>AP</u>	+!		<u>180MG/20ML (9MG/ML)</u>	<u>N018803 002</u> Oct 29, 1982
<u>AP</u>	+!		<u>450MG/50ML (9MG/ML)</u>	<u>N018803 003</u> Oct 29, 1982
<u>AP</u>	+		<u>450MG/50ML (9MG/ML)</u>	<u>N019465 002</u> Jul 15, 1985
<u>AP</u>	+!		<u>900MG/100ML</u>	<u>N019465 001</u> Jul 15, 1985
<u>AP</u>		NEPHRON	<u>450MG/50ML (9MG/ML)</u>	<u>A211968 001</u> Apr 23, 2020
<u>AP</u>			<u>900MG/100ML</u>	<u>A211968 002</u> Apr 23, 2020
<u>AP</u>	!	TARO	<u>45MG/5ML (9MG/ML)</u>	<u>A077407 001</u> Aug 11, 2006
<u>AP</u>	!		<u>90MG/10ML (9MG/ML)</u>	<u>A077407 002</u> Aug 11, 2006

SODIUM CHLORIDE 0.9%

+	B BRAUN	90MG/10ML (9MG/ML)	N019635 005	Aug 08, 2016
	HIKMA	18MG/2ML (9MG/ML)	A201833 001	Sep 24, 2013

SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

+!	BAXTER HLTHCARE	225MG/25ML (9MG/ML)	N016677 004	Oct 30, 1985
!	FRESENIUS KABI USA	18MG/2ML (9MG/ML)	A088912 004	Nov 29, 1985

SOLUTION; INTRAVENOUS

SODIUM CHLORIDE 14.6%

<u>AP</u>	FRESENIUS KABI USA	<u>100MEQ/40ML (2.5MEQ/ML)</u>	<u>A212070 002</u>	Apr 28, 2021
<u>AP</u>	+!	HOSPIRA	<u>100MEQ/40ML (2.5MEQ/ML)</u>	<u>N018897 002</u> Jul 20, 1984

SODIUM CHLORIDE 23.4%

<u>AP</u>	FRESENIUS KABI USA	<u>400MEQ/100ML (4MEQ/ML)</u>	<u>A212070 003</u>	Feb 14, 2022
<u>AP</u>	+!	HOSPIRA	<u>400MEQ/100ML (4MEQ/ML)</u>	<u>N018897 003</u> Jun 18, 2020

SODIUM CHLORIDE 14.6%

	FRESENIUS KABI USA	50MEQ/20ML (2.5MEQ/ML)	A212070 001	Apr 28, 2021
--	--------------------	------------------------	-------------	--------------

SOLUTION; IRRIGATION

SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

<u>AT</u>	+!	B BRAUN	<u>900MG/100ML</u>	<u>N016733 001</u>
<u>AT</u>		BAXTER HLTHCARE	<u>900MG/100ML</u>	<u>N017427 001</u>
<u>AT</u>			<u>900MG/100ML</u>	<u>N017867 001</u>
<u>AT</u>		FRESENIUS KABI USA	<u>900MG/100ML</u>	<u>A213688 001</u> Jun 24, 2024
<u>AT</u>	+	ICU MEDICAL INC	<u>900MG/100ML</u>	<u>N017514 001</u>
<u>AT</u>			<u>900MG/100ML</u>	<u>N018314 001</u>

SOLUTION FOR SLUSH; IRRIGATION

SODIUM CHLORIDE 0.9% IN STERILE PLASTIC CONTAINER

+!	BAXTER HLTHCARE	900MG/100ML	N019319 002	May 17, 1985
----	-----------------	-------------	-------------	--------------

SODIUM FLUORIDE F-18

INJECTABLE; INTRAVENOUS

SODIUM FLUORIDE F-18

<u>AP</u>		3D IMAGING DRUG	<u>10-200mCi/ML</u>	<u>A203777 001</u> Oct 19, 2015
<u>AP</u>		BAMF	<u>10-200mCi/ML</u>	<u>A216126 001</u> Dec 06, 2022
<u>AP</u>		BIOMEDCL RES FDN	<u>10-200mCi/ML</u>	<u>A204351 001</u> Jan 09, 2015
<u>AP</u>		CARDINAL HEALTH 414	<u>10-200mCi/ML</u>	<u>A203780 001</u> Jul 30, 2015
<u>AP</u>		ESSENTIAL ISOTOPES	<u>10-200mCi/ML</u>	<u>A204541 001</u> Oct 29, 2014
<u>AP</u>		HOT SHOTS NM LLC	<u>10-200mCi/ML</u>	<u>A204530 001</u> Jul 29, 2015
<u>AP</u>		JUBILANT DRAXIMAGE	<u>10-200mCi/ML</u>	<u>A203968 001</u> Oct 23, 2015
<u>AP</u>		KREITCHMAN PET CTR	<u>10-200mCi/ML</u>	<u>A203936 001</u> May 19, 2016
<u>AP</u>		MIDWEST MEDCL	<u>10-200mCi/ML</u>	<u>A204440 001</u> Nov 17, 2015
<u>AP</u>		MIPS CRF	<u>10-200mCi/ML</u>	<u>A204517 001</u> Jul 21, 2015
<u>AP</u>		NCM USA BRONX LLC	<u>10-200mCi/ML</u>	<u>A204513 001</u> Nov 28, 2014
<u>AP</u>		NUKEMED	<u>10-200mCi/ML</u>	<u>A203912 001</u> Apr 22, 2015
<u>AP</u>	!	PETNET	<u>10-200mCi/ML</u>	<u>A203890 001</u> Sep 28, 2015
<u>AP</u>		PRECISION NUCLEAR	<u>10-200mCi/ML</u>	<u>A204542 001</u> Feb 27, 2015
<u>AP</u>		SOFIE	<u>10-200mCi/ML</u>	<u>A203592 001</u> Aug 18, 2015
<u>AP</u>		UNIV UTAH CYCLOTRON	<u>10-200mCi/ML</u>	<u>A204497 001</u> Apr 20, 2015
	!	MCPRF	10-91.5mCi/ML	A203605 001 Jun 28, 2013
		THE FEINSTEIN INST	20-600mCi/ML	A204328 001 Nov 19, 2014

PRESCRIPTION DRUG PRODUCT LIST

SODIUM IODIDE I-123

CAPSULE; ORAL

SODIUM IODIDE I 123

AA	+ !	CARDINAL HEALTH 418	100uCi	N018671 001	May 27, 1982
AA	+ !		200uCi	N018671 002	May 27, 1982
AA		CURIUM	100uCi	A071909 001	Feb 28, 1989
AA			200uCi	A071910 001	Feb 28, 1989

SODIUM IODIDE I-131

CAPSULE; ORAL

SODIUM IODIDE I 131

+ JUBILANT

0.009-0.1mCi

N021305 006 May 19, 2005

SOLUTION; ORAL

HICON

AA	+ !	JUBILANT	250-1000mCi	N021305 007	Dec 05, 2011
AA		INTL ISOTOPES	250-1000mCi	A209166 001	Feb 05, 2020

SODIUM NITRITE

SOLUTION; INTRAVENOUS

SODIUM NITRITE

+! HOPE PHARMS

300MG/10ML (30MG/ML)

N203922 001 Feb 14, 2012

SODIUM NITRITE; SODIUM THIOSULFATE

SOLUTION, SOLUTION; INTRAVENOUS, INTRAVENOUS

NITHIODET

+! HOPE PHARMS

300MG/10ML (30MG/ML), N/A; N/A, 12.5GM/50ML
(250MG/ML)

N201444 001 Jan 14, 2011

SODIUM NITROPRUSSIDE

INJECTABLE; INJECTION

SODIUM NITROPRUSSIDE

AP		AMNEAL	25MG/ML	A209493 001	Nov 07, 2017
AP		BE PHARMS	25MG/ML	A214971 001	Jul 12, 2021
AP		CAPLIN	25MG/ML	A211016 001	Nov 29, 2019
AP		DR REDDYS	25MG/ML	A210114 001	Apr 10, 2019
AP	!	MICRO LABS	25MG/ML	A209352 001	Dec 08, 2017
AP		MYLAN LABS LTD	25MG/ML	A210763 001	Apr 17, 2018
AP		NEXUS	25MG/ML	A207499 001	May 25, 2017
AP		NORVIUM BIOSCIENCE	25MG/ML	A209584 001	Aug 10, 2018
AP		SAGENT PHARMS INC	25MG/ML	A207426 001	Dec 08, 2016
AP		SLATE RUN PHARMA	25MG/ML	A214199 001	Aug 25, 2020
AP		SOMERSET THERAPS LLC	25MG/ML	A210882 001	Aug 17, 2018
AP		XIROMED	25MG/ML	A211277 001	Oct 29, 2020
		SOLUTION; INTRAVENOUS			
		<u>NIPRIDE RTU IN SODIUM CHLORIDE 0.9%</u>			
AP	+ !	EXELA PHARMA	20MG/100ML (0.2MG/ML)	N209387 003	Jul 13, 2018
AP	+ !		50MG/100ML (0.5MG/ML)	N209387 001	Mar 08, 2017
		<u>SODIUM NITROPRUSSIDE</u>			
AP		SLATE RUN PHARMA	20MG/100ML (0.2MG/ML)	A215846 002	Aug 26, 2022
AP			50MG/100ML (0.5MG/ML)	A215846 003	Aug 26, 2022
			10MG/50ML (0.2MG/ML)	A215846 001	Aug 26, 2022

SODIUM OXYBATE

FOR SUSPENSION, EXTENDED RELEASE; ORAL

LUMRYZ

+! AVADEL CNS

4.5GM/PACKET

N214755 001 May 01, 2023

+

6GM/PACKET

N214755 002 May 01, 2023

+

7.5GM/PACKET

N214755 003 May 01, 2023

+

9GM/PACKET

N214755 004 May 01, 2023

SOLUTION; ORAL

XYREM

+! JAZZ PHARMS

0.5GM/ML

N021196 001 Jul 17, 2002

SODIUM PHENYLBUTYRATE

FOR SUSPENSION; ORAL

OLPRUVA

+ ACER

2GM/PACKET

N214860 001 Dec 22, 2022

+

3GM/PACKET

N214860 002 Dec 22, 2022

+

4GM/PACKET

N214860 003 Dec 22, 2022

+!

5GM/PACKET

N214860 004 Dec 22, 2022

+

6GM/PACKET

N214860 005 Dec 22, 2022

+

6.67GM/PACKET

N214860 006 Dec 22, 2022

PRESCRIPTION DRUG PRODUCT LIST

SODIUM PHENYLBUTYRATE

PELLETS; ORAL

PHEBURANE

+! MEDUNIK

84GM/BOT

N216513 001 Jun 17, 2022

POWDER; ORAL

BUPHENYL**AB** +! HORIZON THERAP US3GM/TEASPOONFULN020573 001 Apr 30, 1996SODIUM PHENYLBUTYRATE**AB** ENDO OPERATIONS3GM/TEASPOONFULA203918 001 Jun 15, 2016**AB** SIGMAPHARM LABS LLC3GM/TEASPOONFULA202819 001 Mar 22, 2013

TABLET; ORAL

BUPHENYL**AB** +! HORIZON THERAP US500MGN020572 001 May 13, 1996SODIUM PHENYLBUTYRATE**AB** ENDO OPERATIONS500MGA204395 001 Apr 15, 2016**AB** GLENMARK PHARMS LTD500MGA216462 001 Nov 01, 2022SODIUM PHOSPHATE, DIBASIC, ANHYDROUS; SODIUM PHOSPHATE, MONOBASIC, MONOHYDRATE

TABLET; ORAL

MONOBASIC SODIUM PHOSPHATE AND DIBASIC SODIUM PHOSPHATE

! NOVEL LABS INC

0.398GM;1.102GM

A079247 001 Dec 30, 2011

SODIUM PHOSPHATE, DIBASIC, HEPTAHYDRATE; SODIUM PHOSPHATE, MONOBASIC, MONOHYDRATE

SOLUTION; INTRAVENOUS

SODIUM PHOSPHATES**AP** AM REGENT1.34GM/5ML;1.38GM/5MLA218314 001 Jun 21, 2024(2.68MG/ML;2.76MG/ML)**AP**4.02GM/15ML;4.14GM/15MLA218314 002 Jun 21, 2024(2.68MG/ML;2.76MG/ML)**AP**13.4GM/50ML;13.8GM/50MLA218314 003 Jun 21, 2024(2.68MG/ML;2.76MG/ML)**AP** ! FRESENIUS KABI USA1.34GM/5ML;1.38GM/5MLA209997 001 Mar 30, 2022(2.68MG/ML;2.76MG/ML)**AP**4.02GM/15ML;4.14GM/15MLA209997 002 Mar 30, 2022(2.68MG/ML;2.76MG/ML)**AP** !13.4GM/50ML;13.8GM/50MLA209997 003 Mar 30, 2022(2.68MG/ML;2.76MG/ML)SODIUM PHOSPHATES IN PLASTIC CONTAINER**AP** +! HOSPIRA4.02GM/15ML;4.14GM/15MLN018892 001 May 10, 1983(2.68MG/ML;2.76MG/ML)SODIUM POLYSTYRENE SULFONATE

POWDER; ORAL, RECTAL

KALEXATE**AA** ! KVK TECH454GM/BOTA040905 001 Mar 30, 2009KIONEX**AA** ANI PHARMS454GM/BOTA040029 001 Feb 06, 1998SODIUM POLYSTYRENE SULFONATE**AA** BELCHER454GM/BOTA205727 001 Feb 23, 2016**AA** CHARTWELL RX454GM/BOTA206815 001 Feb 18, 2016**AA** CMP PHARMA INC454GM/BOTA089910 001 Jan 19, 1989**AA** EPIC PHARMA LLC453.6GM/BOTA202333 001 Mar 19, 2014**AA** NUVO PHARMS INC454GM/BOTA204071 001 Nov 28, 2014**AA** UPSHER SMITH LABS453.6GM/BOTA090313 001 Dec 21, 2011

KALEXATE

KVK TECH

15GM/BOT

A040905 002 Apr 03, 2015

SODIUM POLYSTYRENE SULFONATE

NUVO PHARMS INC

15GM/BOT

A204071 002 Nov 28, 2014

SUSPENSION; ORAL, RECTAL

KIONEX**AA** ANI PHARMS15GM/60MLA040028 001 Sep 17, 2007SPS**AA** +! CMP PHARMA INC15GM/60MLA087859 001 Dec 08, 1982SODIUM TETRADECYL SULFATE

INJECTABLE; INJECTION

SODIUM TETRADECYL SULFATE**AP** HIKMA60MG/2ML (30MG/ML)A209937 001 Dec 09, 2019SOTRADECOL**AP** ! MYLAN INSTITUTIONAL60MG/2ML (30MG/ML)A040541 002 Nov 12, 2004

!

20MG/2ML (10MG/ML)

A040541 001 Nov 12, 2004

PRESCRIPTION DRUG PRODUCT LISTSODIUM THIOSULFATE

SOLUTION; INTRAVENOUS

PEDMARK

+! FENNEC PHARMS INC 12.5GM/100ML (125MG/ML) N212937 001 Sep 20, 2022

SODIUM THIOSULFATE

+! HOPE PHARMS 12.5GM/50ML (250MG/ML) N203923 001 Feb 14, 2012

SODIUM ZIRCONIUM CYCLOSILICATE

FOR SUSPENSION; ORAL

LOKELMA

+ ASTRAZENECA 5GM/PACKET N207078 001 May 18, 2018

+! 10GM/PACKET N207078 002 May 18, 2018

SOFOBUVIR

PELLETS; ORAL

SOVALDI

+ GILEAD SCIENCES INC 150MG/PACKET N212480 001 Aug 28, 2019

+! 200MG/PACKET N212480 002 Aug 28, 2019

TABLET; ORAL

SOVALDI

+ GILEAD SCIENCES INC 200MG N204671 002 Aug 28, 2019

+! 400MG N204671 001 Dec 06, 2013

SOFOBUVIR; VELPATASVIR

PELLETS; ORAL

EPCLUSA

+ GILEAD SCIENCES INC 150MG; 37.5MG/PACKET N214187 001 Jun 10, 2021

+! 200MG; 50MG/PACKET N214187 002 Jun 10, 2021

TABLET; ORAL

EPCLUSA

+ GILEAD SCIENCES INC 200MG; 50MG N208341 002 Mar 19, 2020

+! 400MG; 100MG N208341 001 Jun 28, 2016

SOFOBUVIR; VELPATASVIR; VOXILAPREVIR

TABLET; ORAL

VOSEVI

+! GILEAD SCIENCES INC 400MG; 100MG; 100MG N209195 001 Jul 18, 2017

SOPPIRONIUM BROMIDE

GEL, METERED; TOPICAL

SOFDRA

+! BOTANIX SB EQ 12.45% BASE (EQ 72MG BASE/ACTUATION) N217347 001 Jun 18, 2024

SOLIFENACIN SUCCINATE

SUSPENSION; ORAL

VESICARE LS

+! ASTELLAS 1MG/ML N209529 001 May 26, 2020

TABLET; ORAL

SOLIFENACIN SUCCINATE

AB	ALEMBIC	5MG	A205575 001	May 20, 2019
AB		10MG	A205575 002	May 20, 2019
AB	ALKEM LABS LTD	5MG	A210224 001	May 20, 2019
AB		10MG	A210224 002	May 20, 2019
AB	ANNORA PHARMA	5MG	A215761 001	Jun 15, 2022
AB		10MG	A215761 002	Jun 15, 2022
AB	AUROBINDO PHARMA	5MG	A206817 001	Dec 27, 2022
AB		10MG	A206817 002	Dec 27, 2022
AB	AUSTARPHARMA	5MG	A210281 001	May 20, 2019
AB		10MG	A210281 002	May 20, 2019
AB	CHARTWELL RX	5MG	A210582 001	May 20, 2019
AB		10MG	A210582 002	May 20, 2019
AB	ESJAY PHARMA	5MG	A212214 001	Sep 26, 2019
AB		10MG	A212214 002	Sep 26, 2019
AB	GLENMARK SPECLT	5MG	A209239 001	May 20, 2019
AB		10MG	A209239 002	May 20, 2019
AB	JUBILANT GENERICS	5MG	A205484 001	Oct 02, 2024
AB		10MG	A205484 002	Oct 02, 2024
AB	MACLEODS PHARMS LTD	5MG	A204374 001	Feb 20, 2024
AB		10MG	A204374 002	Feb 20, 2024
AB	MSN	5MG	A210688 001	May 20, 2019
AB		10MG	A210688 002	May 20, 2019
AB	QILU	5MG	A209333 001	May 20, 2019
AB		10MG	A209333 002	May 20, 2019
AB	RISING	5MG	A211423 001	Dec 11, 2019
AB		10MG	A211423 002	Dec 11, 2019

PRESCRIPTION DRUG PRODUCT LIST

SOLIFENACIN SUCCINATE

TABLET;ORAL

SOLIFENACIN SUCCINATE

<u>AB</u>	SCIEGEN PHARMS INC	<u>5MG</u>	<u>A211657</u>	<u>001</u>	May 20, 2019
<u>AB</u>		<u>10MG</u>	<u>A211657</u>	<u>002</u>	May 20, 2019
<u>AB</u>	TEVA PHARMS USA	<u>5MG</u>	<u>A091464</u>	<u>001</u>	Apr 02, 2014
<u>AB</u>		<u>10MG</u>	<u>A091464</u>	<u>002</u>	Apr 02, 2014
<u>AB</u>	UNICHEM	<u>5MG</u>	<u>A211701</u>	<u>001</u>	Aug 27, 2019
<u>AB</u>		<u>10MG</u>	<u>A211701</u>	<u>002</u>	Aug 27, 2019
<u>AB</u>	WATSON LABS INC	<u>5MG</u>	<u>A202551</u>	<u>001</u>	May 20, 2019
<u>AB</u>		<u>10MG</u>	<u>A202551</u>	<u>002</u>	May 20, 2019

VESICARE

<u>AB</u>	+ ASTELLAS	<u>5MG</u>	<u>N021518</u>	<u>001</u>	Nov 19, 2004
<u>AB</u>	+!	<u>10MG</u>	<u>N021518</u>	<u>002</u>	Nov 19, 2004

SOLRIAMFETOL HYDROCHLORIDE

TABLET;ORAL

SUNOSI

+	AXSOME MALTA	EQ 75MG BASE	N211230	001	Jun 17, 2019
+	!	EQ 150MG BASE	N211230	002	Jun 17, 2019

SONIDEGIB PHOSPHATE

CAPSULE;ORAL

ODOMZO

+	SUN PHARM	EQ 200MG BASE	N205266	001	Jul 24, 2015
---	-----------	---------------	---------	-----	--------------

SORAFENIB TOSYLATE

TABLET;ORAL

NEXAVAR

<u>AB</u>	+! BAYER HLTHCARE	<u>EQ 200MG BASE</u>	<u>N021923</u>	<u>001</u>	Dec 20, 2005
-----------	-------------------	----------------------	----------------	------------	--------------

SORAFENIB TOSYLATE

<u>AB</u>	DR REDDYS	<u>EQ 200MG BASE</u>	<u>A216073</u>	<u>001</u>	Jun 07, 2022
<u>AB</u>	MYLAN	<u>EQ 200MG BASE</u>	<u>A207012</u>	<u>001</u>	Sep 10, 2020
<u>AB</u>	TEVA PHARMS USA INC	<u>EQ 200MG BASE</u>	<u>A209567</u>	<u>001</u>	Nov 12, 2020
<u>AB</u>	TORRENT	<u>EQ 200MG BASE</u>	<u>A217095</u>	<u>001</u>	Apr 12, 2023
<u>AB</u>	YABAO PHARM	<u>EQ 200MG BASE</u>	<u>A209050</u>	<u>001</u>	Nov 09, 2022

SORBITOL

SOLUTION;IRRIGATION

SORBITOL 3% IN PLASTIC CONTAINER

	BAXTER HLTHCARE	3GM/100ML	N017863	001	
--	-----------------	-----------	---------	-----	--

SORBITOL 3.3% IN PLASTIC CONTAINER

	B BRAUN	3.3GM/100ML	N016741	001	
--	---------	-------------	---------	-----	--

SOTAGLIFLOZIN

TABLET;ORAL

INPEFA

+	LEXICON PHARMS INC	200MG	N216203	001	May 26, 2023
+	!	400MG	N216203	002	May 26, 2023

SOTALOL HYDROCHLORIDE

SOLUTION;INTRAVENOUS

SOTALOL HYDROCHLORIDE

+	! ALTATHERA PHARMS LLC	150MG/10ML (15MG/ML)	N022306	001	Jul 02, 2009
---	------------------------	----------------------	---------	-----	--------------

SOLUTION;ORAL

SOTYLIZE

+	! AZURITY	5MG/ML (5MG/ML)	N205108	001	Oct 22, 2014
---	-----------	-----------------	---------	-----	--------------

TABLET;ORAL

BETAPACE

<u>AB</u>	+ LEGACY PHARMA	<u>80MG</u>	<u>N019865</u>	<u>001</u>	Oct 30, 1992
<u>AB</u>	+	<u>120MG</u>	<u>N019865</u>	<u>005</u>	Apr 20, 1994
<u>AB</u>	+!	<u>160MG</u>	<u>N019865</u>	<u>002</u>	Oct 30, 1992
<u>AB</u>	+	<u>240MG</u>	<u>N019865</u>	<u>003</u>	Oct 30, 1992

BETAPACE AF

<u>AB</u>	+ LEGACY PHARMA	<u>80MG</u>	<u>N021151</u>	<u>001</u>	Feb 22, 2000
<u>AB</u>	+	<u>120MG</u>	<u>N021151</u>	<u>002</u>	Feb 22, 2000
<u>AB</u>	+!	<u>160MG</u>	<u>N021151</u>	<u>003</u>	Feb 22, 2000

SORINE

<u>AB</u>	AIPING PHARM INC	<u>80MG</u>	<u>A075500</u>	<u>001</u>	Apr 27, 2001
<u>AB</u>		<u>120MG</u>	<u>A075500</u>	<u>004</u>	Apr 27, 2001
<u>AB</u>		<u>160MG</u>	<u>A075500</u>	<u>002</u>	Apr 27, 2001
<u>AB</u>		<u>240MG</u>	<u>A075500</u>	<u>003</u>	Apr 27, 2001

SOTALOL HYDROCHLORIDE

<u>AB</u>	APOTEX	<u>80MG</u>	<u>A076140</u>	<u>001</u>	Sep 26, 2002
<u>AB</u>		<u>80MG</u>	<u>A076214</u>	<u>001</u>	Aug 27, 2003

PRESCRIPTION DRUG PRODUCT LIST

SOTALOL HYDROCHLORIDE

TABLET; ORAL

SOTALOL HYDROCHLORIDE

<u>AB</u>		<u>120MG</u>	<u>A076140 002</u>	Sep 26, 2002
<u>AB</u>		<u>120MG</u>	<u>A076214 002</u>	Aug 27, 2003
<u>AB</u>		<u>160MG</u>	<u>A076140 003</u>	Sep 26, 2002
<u>AB</u>		<u>160MG</u>	<u>A076214 003</u>	Aug 27, 2003
<u>AB</u>		<u>240MG</u>	<u>A076140 004</u>	Sep 26, 2002
<u>AB</u>	AUROBINDO PHARMA USA	<u>80MG</u>	<u>A077616 001</u>	Feb 07, 2007
<u>AB</u>		<u>120MG</u>	<u>A077616 002</u>	Feb 07, 2007
<u>AB</u>		<u>160MG</u>	<u>A077616 003</u>	Feb 07, 2007
<u>AB</u>	BEXIMCO PHARMS USA	<u>80MG</u>	<u>A207428 001</u>	Oct 21, 2016
<u>AB</u>		<u>80MG</u>	<u>A207429 001</u>	Nov 02, 2018
<u>AB</u>		<u>120MG</u>	<u>A207428 002</u>	Oct 21, 2016
<u>AB</u>		<u>120MG</u>	<u>A207429 002</u>	Nov 02, 2018
<u>AB</u>		<u>160MG</u>	<u>A207428 003</u>	Oct 21, 2016
<u>AB</u>		<u>160MG</u>	<u>A207429 003</u>	Nov 02, 2018
<u>AB</u>	EPIC PHARMA LLC	<u>80MG</u>	<u>A077070 001</u>	Nov 04, 2005
<u>AB</u>		<u>120MG</u>	<u>A077070 002</u>	Nov 04, 2005
<u>AB</u>		<u>160MG</u>	<u>A077070 003</u>	Nov 04, 2005
<u>AB</u>	OXFORD PHARMS	<u>80MG</u>	<u>A075563 001</u>	Nov 07, 2003
<u>AB</u>		<u>120MG</u>	<u>A075563 002</u>	Nov 07, 2003
<u>AB</u>		<u>160MG</u>	<u>A075563 003</u>	Nov 07, 2003
<u>AB</u>		<u>240MG</u>	<u>A075563 004</u>	Nov 07, 2003
<u>AB</u>	TEVA	<u>80MG</u>	<u>A075429 001</u>	May 01, 2000
<u>AB</u>		<u>120MG</u>	<u>A075429 002</u>	May 01, 2000
<u>AB</u>		<u>160MG</u>	<u>A075429 003</u>	May 01, 2000
<u>AB</u>		<u>240MG</u>	<u>A075429 004</u>	May 01, 2000

SOTORASIB

TABLET; ORAL

LUMAKRAS

+	AMGEN INC	120MG	N214665 001	May 28, 2021
+		240MG	N214665 003	Jun 26, 2024
+	!	320MG	N214665 002	Jan 20, 2023

SOYBEAN OIL

INJECTABLE; INJECTION

INTRALIPID 20%

<u>AP</u>	+	FRESENIUS	<u>20%</u>	<u>N018449 001</u>	
<u>AP</u>	+	!	<u>20%</u>	<u>N020248 001</u>	Aug 07, 1996
			<u>NUTRILIPID 20%</u>		
<u>AP</u>	+	B BRAUN	<u>20%</u>	<u>N019531 002</u>	May 28, 1993
			<u>INTRALIPID 30%</u>		
	+	FRESENIUS	30%	N019942 001	Dec 30, 1993

SPARSENTAN

TABLET; ORAL

FILSPARI

+	TRAVERE	200MG	N216403 001	Feb 17, 2023
+	!	400MG	N216403 002	Feb 17, 2023

SPINOSAD

SUSPENSION; TOPICAL

NATROBA

+	CIPHER	0.9%	N022408 001	Jan 18, 2011
---	--------	------	-------------	--------------

SPIRONOLACTONE

SUSPENSION; ORAL

CAROSPIR

<u>AB</u>	+	CMP DEV LLC	<u>25MG/5ML</u>	<u>N209478 001</u>	Aug 04, 2017
-----------	---	-------------	-----------------	--------------------	--------------

SPIRONOLACTONE

<u>AB</u>		AMNEAL	<u>25MG/5ML</u>	<u>A215572 001</u>	Sep 05, 2023
-----------	--	--------	-----------------	--------------------	--------------

TABLET; ORAL

ALDACTONE

<u>AB</u>	+	PFIZER	<u>25MG</u>	<u>N012151 009</u>	Dec 30, 1983
<u>AB</u>	+		<u>50MG</u>	<u>N012151 008</u>	Dec 30, 1982
<u>AB</u>	+	!	<u>100MG</u>	<u>N012151 010</u>	Dec 30, 1983

SPIRONOLACTONE

<u>AB</u>		ACCORD HLTHCARE	<u>25MG</u>	<u>A203512 001</u>	Sep 19, 2016
<u>AB</u>			<u>50MG</u>	<u>A203512 002</u>	Sep 19, 2016
<u>AB</u>			<u>100MG</u>	<u>A203512 003</u>	Sep 19, 2016
<u>AB</u>		AMNEAL PHARMS	<u>25MG</u>	<u>A091426 001</u>	Jul 02, 2010
<u>AB</u>			<u>50MG</u>	<u>A091426 002</u>	Jul 02, 2010

PRESCRIPTION DRUG PRODUCT LIST

SPIRONOLACTONE

TABLET;ORAL

SPIRONOLACTONE

<u>AB</u>		<u>100MG</u>	<u>A091426</u>	<u>003</u>	Jul 02, 2010
<u>AB</u>	AUROBINDO PHARMA	<u>25MG</u>	<u>A202187</u>	<u>001</u>	Mar 06, 2014
<u>AB</u>		<u>50MG</u>	<u>A202187</u>	<u>002</u>	Mar 06, 2014
<u>AB</u>		<u>100MG</u>	<u>A202187</u>	<u>003</u>	Mar 06, 2014
<u>AB</u>	JUBILANT GENERICS	<u>25MG</u>	<u>A203253</u>	<u>001</u>	Apr 23, 2014
<u>AB</u>		<u>50MG</u>	<u>A203253</u>	<u>002</u>	Apr 23, 2014
<u>AB</u>		<u>100MG</u>	<u>A203253</u>	<u>003</u>	Apr 23, 2014
<u>AB</u>	MYLAN	<u>25MG</u>	<u>A040424</u>	<u>001</u>	Aug 20, 2001
<u>AB</u>		<u>50MG</u>	<u>A040424</u>	<u>002</u>	Aug 20, 2001
<u>AB</u>		<u>100MG</u>	<u>A040424</u>	<u>003</u>	Aug 20, 2001
<u>AB</u>	OXFORD PHARMS	<u>25MG</u>	<u>A040750</u>	<u>001</u>	Aug 29, 2006
<u>AB</u>		<u>50MG</u>	<u>A040750</u>	<u>002</u>	Aug 29, 2006
<u>AB</u>		<u>100MG</u>	<u>A040750</u>	<u>003</u>	Aug 29, 2006
<u>AB</u>	SUN PHARM INDUSTRIES	<u>25MG</u>	<u>A089424</u>	<u>001</u>	Jul 23, 1986
<u>AB</u>		<u>50MG</u>	<u>A089424</u>	<u>002</u>	Aug 11, 1999
<u>AB</u>		<u>100MG</u>	<u>A089424</u>	<u>003</u>	Aug 11, 1999
<u>AB</u>	ZYDUS PHARMS	<u>25MG</u>	<u>A205936</u>	<u>001</u>	Jul 18, 2018
<u>AB</u>		<u>50MG</u>	<u>A205936</u>	<u>002</u>	Jul 18, 2018
<u>AB</u>		<u>100MG</u>	<u>A205936</u>	<u>003</u>	Jul 18, 2018

STERILE WATER FOR INJECTION

LIQUID;N/A

BACTERIOSTATIC WATER FOR INJECTION IN PLASTIC CONTAINER

<u>AP</u>	<u>+</u> !	HOSPIRA	<u>100%</u>	<u>N018802</u>	<u>001</u>	Oct 27, 1982
-----------	------------	---------	-------------	----------------	------------	--------------

STERILE WATER FOR INJECTION

<u>AP</u>		AM REGENT	<u>100% (10ML)</u>	<u>A217341</u>	<u>001</u>	Aug 29, 2023
<u>AP</u>			<u>100% (20ML)</u>	<u>A217341</u>	<u>002</u>	Aug 29, 2023
<u>AP</u>			<u>100% (5ML)</u>	<u>A217341</u>	<u>003</u>	Feb 09, 2024
<u>AP</u>		FRESENIUS KABI USA	<u>100%</u>	<u>A209689</u>	<u>001</u>	Nov 24, 2017
<u>AP</u>		HIKMA	<u>100% (10ML)</u>	<u>A206369</u>	<u>001</u>	Sep 02, 2015
<u>AP</u>			<u>100%</u>	<u>A212735</u>	<u>001</u>	Jan 31, 2023
<u>AP</u>	<u>+</u> !	HOSPIRA	<u>100% (10ML)</u>	<u>N018801</u>	<u>002</u>	Oct 27, 1982
<u>AP</u>	<u>+</u> !		<u>100% (20ML)</u>	<u>N018801</u>	<u>003</u>	Oct 27, 1982
<u>AP</u>	<u>!</u>	MEDEFIL INC	<u>100% (5ML)</u>	<u>A211188</u>	<u>004</u>	Dec 02, 2019
<u>AP</u>			<u>100% (10ML)</u>	<u>A211188</u>	<u>005</u>	Dec 02, 2019
<u>AP</u>		NEPHRON	<u>100% (5ML)</u>	<u>A211222</u>	<u>001</u>	Feb 10, 2021
<u>AP</u>		NEXUS	<u>100% (10ML)</u>	<u>A217536</u>	<u>001</u>	Jul 11, 2023
<u>AP</u>			<u>100% (20ML)</u>	<u>A217536</u>	<u>002</u>	Jul 11, 2023

STERILE WATER FOR INJECTION IN PLASTIC CONTAINER

<u>AP</u>	<u>+</u> !	B BRAUN	<u>100%</u>	<u>N019633</u>	<u>001</u>	Feb 29, 1988
<u>AP</u>	<u>+</u> !	BAXTER HLTHCARE	<u>100%</u>	<u>N018632</u>	<u>001</u>	Jun 30, 1982
<u>AP</u>	<u>+</u> !		<u>100%</u>	<u>N018632</u>	<u>002</u>	Apr 19, 1988
<u>AP</u>		FRESENIUS KABI USA	<u>100%</u>	<u>A088400</u>	<u>001</u>	Jan 16, 1984
<u>AP</u>	<u>+</u> !	ICU MEDICAL INC	<u>100%</u>	<u>N018233</u>	<u>001</u>	
<u>AP</u>	<u>+</u> !		<u>100%</u>	<u>N019869</u>	<u>001</u>	Dec 26, 1989
<u>AP</u>		TARO	<u>100%</u>	<u>A077393</u>	<u>001</u>	Aug 11, 2006

STERILE WATER FOR INJECTION

<u>+</u> !	HOSPIRA	100% (50ML)	N018801	004	Oct 27, 1982
<u>+</u> !		100% (100ML)	N018801	005	Oct 27, 1982
	MEDEFIL INC	100% (1ML)	A211188	001	Dec 02, 2019
<u>!</u>		100% (2.5ML)	A211188	002	Dec 02, 2019
<u>!</u>		100% (3ML)	A211188	003	Dec 02, 2019

STERILE WATER FOR IRRIGATION

LIQUID;IRRIGATION

STERILE WATER

<u>AT</u>	<u>+</u>	BAXTER HLTHCARE	<u>100%</u>	<u>N017428</u>	<u>001</u>
-----------	----------	-----------------	-------------	----------------	------------

STERILE WATER FOR IRRIGATION

<u>AT</u>		FRESENIUS KABI USA	<u>100%</u>	<u>A216123</u>	<u>001</u>	Aug 12, 2024
-----------	--	--------------------	-------------	----------------	------------	--------------

STERILE WATER IN PLASTIC CONTAINER

<u>AT</u>	<u>+</u>	B BRAUN	<u>100%</u>	<u>N016734</u>	<u>001</u>
<u>AT</u>		BAXTER HLTHCARE	<u>100%</u>	<u>N017866</u>	<u>001</u>
<u>AT</u>		ICU MEDICAL INC	<u>100%</u>	<u>N017513</u>	<u>001</u>
<u>AT</u>			<u>100%</u>	<u>N018313</u>	<u>001</u>

PRESCRIPTION DRUG PRODUCT LIST

STIRIPENTOL

CAPSULE; ORAL

DIACOMIT

+ BIOCDEX SA

250MG

N206709 001 Aug 20, 2018

+!

500MG

N206709 002 Aug 20, 2018

FOR SUSPENSION; ORAL

DIACOMIT

+ BIOCDEX SA

250MG/PACKET

N207223 001 Aug 20, 2018

+!

500MG/PACKET

N207223 002 Aug 20, 2018

STREPTOMYCIN SULFATE

INJECTABLE; INJECTION

STREPTOMYCIN SULFATE

! XGEN PHARMS

EQ 1GM BASE/VIAL

A064210 001 Jun 30, 1998

STREPTOZOCIN

INJECTABLE; INJECTION

ZANOSAR

+! TEVA PHARMS USA

1GM/VIAL

N050577 001 May 07, 1982

STRONTIUM CHLORIDE SR-89

INJECTABLE; INJECTION

METASTRON**AP** +! Q BIOMED**1mCi/ML****N020134 001** Jun 18, 1993STRONTIUM CHLORIDE SR-89**AP** Q BIOMED**1mCi/ML****A075941 001** Jan 06, 2003SUCCIMER

CAPSULE; ORAL

CHEMET

+! RECORDATI RARE

100MG

N019998 002 Jan 30, 1991

SUCCINYLCHOLINE CHLORIDE

INJECTABLE; INJECTION

ANECTINE**AP** +! SANDOZ**20MG/ML****N008453 002**QUELICIN**AP** +! HOSPIRA**20MG/ML****N008845 006**SUCCINYLCHOLINE CHLORIDE**AP** ADAPTIS**20MG/ML****A214308 001** May 22, 2020**AP** AMNEAL**20MG/ML****A211432 001** Nov 16, 2018**AP** AMRING PHARMS**20MG/ML****A210231 001** Jun 04, 2018**AP** ASPIRO**20MG/ML****A213810 001** May 04, 2020**AP** BAXTER HLTHCARE**20MG/ML****A218878 001** Sep 24, 2024

CORP

AP BE PHARMS**20MG/ML****A216003 001** Feb 07, 2022**AP** DEVA HOLDING AS**20MG/ML****A214491 001** Dec 21, 2020**AP** DR REDDYS**20MG/ML****A210698 001** Aug 02, 2019**AP** FRESENIUS KABI USA**20MG/ML****A211346 001** Nov 20, 2020**AP** GLAND PHARMA LTD**20MG/ML****A214246 001** Jun 16, 2020**AP** HIKMA**20MG/ML****A213229 001** Jun 12, 2020**AP** MEITHEAL**20MG/ML****A214514 001** Oct 19, 2021**AP** MICRO LABS**20MG/ML****A214879 001** Nov 24, 2020**AP** SAGENT PHARMS INC**20MG/ML****A215022 001** Mar 29, 2021**AP** SOMERSET THERAPS**20MG/ML****A211589 001** Jan 15, 2020

LLC

AP UMEDICA**20MG/ML****A211625 001** May 19, 2020**AP** ! ZYDUS PHARMS**20MG/ML****A209467 001** May 04, 2018

SOLUTION; INTRAMUSCULAR, INTRAVENOUS

SUCCINYLCHOLINE CHLORIDE**AP** DR REDDYS**100MG/5ML (20MG/ML)****A218467 001** Jan 17, 2024**AP** +! HIKMA**100MG/5ML (20MG/ML)****N215143 001** Aug 20, 2021**AP** STERISCIENCE**100MG/5ML (20MG/ML)****A217873 001** Jul 09, 2024SUCRALFATE

SUSPENSION; ORAL

CARAFATE**AB** +! ABBVIE**1GM/10ML****N019183 001** Dec 16, 1993SUCRALFATE**AB** ABON PHARMS LLC**1GM/10ML****A216726 001** Mar 21, 2024**AB** AMNEAL**1GM/10ML****A209356 001** Dec 02, 2019**AB** COSETTE**1GM/10ML****A217126 001** Nov 13, 2024**AB** MYLAN**1GM/10ML****A212913 001** Sep 12, 2022**AB** PD PARTNERS**1GM/10ML****A213549 001** Jan 17, 2024**AB** STRIDES PHARMA**1GM/10ML****A216474 001** May 21, 2024**AB** VISTAPHARM LLC**1GM/10ML****A211884 001** Mar 15, 2022

PRESCRIPTION DRUG PRODUCT LIST

SUCRALFATE

TABLET; ORAL

CARAFATE

AB	+ !	ABBVIE	1GM	N018333	001	
-----------	------------	--------	------------	----------------	------------	--

SUCRALFATE

AB		AMNEAL PHARMS	1GM	A215576	001	Apr 15, 2022
AB		NOSTRUM LABS INC	1GM	A074415	001	Jun 08, 1998
AB		TEVA	1GM	A070848	001	Mar 29, 1996
AB		ZYDUS LIFESCIENCES	1GM	A215705	001	May 03, 2023

SUFENTANIL CITRATE

INJECTABLE; INJECTION

SUFENTA PRESERVATIVE FREE

AP	+ !	RISING	EQ 0.05MG BASE/ML	N019050	001	May 04, 1984
-----------	------------	--------	--------------------------	----------------	------------	--------------

SUFENTANIL CITRATE

AP		HIKMA	EQ 0.05MG BASE/ML	A074413	001	Dec 15, 1995
AP		HOSPIRA	EQ 0.05MG BASE/ML	A074534	001	Dec 11, 1996

SUGAMMADEX SODIUM

SOLUTION; INTRAVENOUS

BRIDION

AP	+	MSD SUB MERCK	EQ 200MG BASE/2ML (EQ 100MG BASE/ML)	N022225	002	Dec 15, 2015
AP	+ !		EQ 500MG BASE/5ML (EQ 100MG BASE/ML)	N022225	001	Dec 15, 2015

SUGAMMADEX SODIUM

AP		ASPIRO	EQ 200MG BASE/2ML (EQ 100MG BASE/ML)	A214337	001	Jun 09, 2023
AP			EQ 500MG BASE/5ML (EQ 100MG BASE/ML)	A214337	002	Jun 09, 2023

SULCONAZOLE NITRATE

CREAM; TOPICAL

EXELDERM

+! JOURNEY

1%

N018737 001 Feb 28, 1989

SOLUTION; TOPICAL

EXELDERM

+! JOURNEY

1%

N018738 001 Aug 30, 1985

SULFACETAMIDE SODIUM

LOTION; TOPICAL

KLARON

AB	+ !	BAUSCH	10%	N019931	001	Dec 23, 1996
-----------	------------	--------	------------	----------------	------------	--------------

SULFACETAMIDE SODIUM

AB		FOUGERA PHARMS	10%	A077015	001	Nov 17, 2006
AB		PADAGIS US	10%	A078649	001	Mar 23, 2009
AB		TARO	10%	A078668	001	May 20, 2009

OINTMENT; OPHTHALMIC

SULFACETAMIDE SODIUM

! PADAGIS US

10%

A080029 001

SOLUTION/DROPS; OPHTHALMIC

SULFACETAMIDE SODIUM

AT	!	BAUSCH AND LOMB	10%	A040066	001	Dec 28, 1994
AT		CHARTWELL RX	10%	A089560	001	Oct 18, 1988

SULFADIAZINE

TABLET; ORAL

SULFADIAZINE

! EPIC PHARMA LLC

500MG

A040091 001 Jul 29, 1994

SULFAMETHOXAZOLE; TRIMETHOPRIM

INJECTABLE; INJECTION

SULFAMETHOXAZOLE AND TRIMETHOPRIM

AP	!	MYLAN LABS LTD	80MG/ML; 16MG/ML	A206607	001	Aug 30, 2017
AP		SOMERSET	80MG/ML; 16MG/ML	A212231	001	Jun 26, 2019
AP		TEVA PHARMS USA	80MG/ML; 16MG/ML	A073303	001	Oct 31, 1991

SUSPENSION; ORAL

SULFAMETHOXAZOLE AND TRIMETHOPRIM

AB	!	AUROBINDO PHARMA	200MG/5ML; 40MG/5ML	A091348	001	Jun 08, 2010
AB		CHARTWELL MOLECULAR	200MG/5ML; 40MG/5ML	A077785	001	Jan 24, 2007
AB		NOVITIUM PHARMA	200MG/5ML; 40MG/5ML	A214330	001	Feb 08, 2022
AB		PRASCO	200MG/5ML; 40MG/5ML	A077612	001	Nov 13, 2006

SULFATRIM PEDIATRIC

AB		PHARM ASSOC	200MG/5ML; 40MG/5ML	N018615	001	Jan 07, 1983
-----------	--	-------------	----------------------------	----------------	------------	--------------

TABLET; ORAL

BACTRIM

AB	+	SUN PHARM INDUSTRIES	400MG; 80MG	N017377	001	
-----------	----------	----------------------	--------------------	----------------	------------	--

PRESCRIPTION DRUG PRODUCT LIST

SULFAMETHOXAZOLE; TRIMETHOPRIM

TABLET; ORAL

BACTRIM DS

AB	+ !	SUN PHARM INDUSTRIES	800MG;160MG	N017377 002	
-----------	------------	-------------------------	--------------------	--------------------	--

SEPTRA

AB		MONARCH PHARMS	400MG;80MG	N017376 001	
-----------	--	----------------	-------------------	--------------------	--

SEPTRA DS

AB		MONARCH PHARMS	800MG;160MG	N017376 002	
-----------	--	----------------	--------------------	--------------------	--

SULFAMETHOXAZOLE AND TRIMETHOPRIM

AB		AMNEAL PHARMS NY	400MG;80MG	A076899 001	Jan 27, 2005
-----------	--	------------------	-------------------	--------------------	--------------

AB			800MG;160MG	A076899 002	Jan 27, 2005
-----------	--	--	--------------------	--------------------	--------------

AB		AUROBINDO PHARMA	400MG;80MG	A090624 001	Feb 16, 2010
-----------	--	------------------	-------------------	--------------------	--------------

AB			800MG;160MG	A090624 002	Feb 16, 2010
-----------	--	--	--------------------	--------------------	--------------

AB		CHARTWELL MOLECULES	400MG;80MG	A078060 002	Jan 25, 2007
-----------	--	---------------------	-------------------	--------------------	--------------

AB			800MG;160MG	A078060 001	Jan 25, 2007
-----------	--	--	--------------------	--------------------	--------------

AB		GLENMARK PHARMS LTD	400MG;80MG	A090828 002	Dec 22, 2010
-----------	--	---------------------	-------------------	--------------------	--------------

AB			800MG;160MG	A090828 001	Dec 22, 2010
-----------	--	--	--------------------	--------------------	--------------

AB		SUN PHARM INDUSTRIES	400MG;80MG	A071017 002	Aug 25, 1986
-----------	--	-------------------------	-------------------	--------------------	--------------

AB			800MG;160MG	A071017 001	Aug 25, 1986
-----------	--	--	--------------------	--------------------	--------------

AB		VISTA PHARMS	400MG;80MG	A076817 001	Oct 07, 2005
-----------	--	--------------	-------------------	--------------------	--------------

AB			800MG;160MG	A076817 002	Oct 07, 2005
-----------	--	--	--------------------	--------------------	--------------

SULFASALAZINE

TABLET; ORAL

AZULFIDINE

AB	+ !	PFIZER	500MG	N007073 001	
-----------	------------	--------	--------------	--------------------	--

SULFASALAZINE

AB		CHARTWELL	500MG	A080197 001	
-----------	--	-----------	--------------	--------------------	--

AB		NUVO PHARMS INC	500MG	A040349 001	Jan 11, 2002
-----------	--	-----------------	--------------	--------------------	--------------

AB		WATSON LABS	500MG	A085828 001	
-----------	--	-------------	--------------	--------------------	--

TABLET, DELAYED RELEASE; ORAL

AZULFIDINE EN-TABS

AB	+ !	PFIZER	500MG	N007073 002	Apr 06, 1983
-----------	------------	--------	--------------	--------------------	--------------

SULFASALAZINE

AB		NUVO PHARMS INC	500MG	A075339 001	Jan 11, 2002
-----------	--	-----------------	--------------	--------------------	--------------

SULFUR HEXAFLUORIDE LIPID-TYPE A MICROSPHERES

FOR SUSPENSION; INTRAVENOUS

LUMASON

+ !	BRACCO	60.7MG/25MG	N203684 001	Oct 15, 2014	
------------	--------	--------------------	--------------------	--------------	--

SULINDAC

TABLET; ORAL

SULINDAC

AB		EPIC PHARMA	150MG	A072710 001	Mar 25, 1991
-----------	--	-------------	--------------	--------------------	--------------

AB			200MG	A072711 001	Mar 25, 1991
-----------	--	--	--------------	--------------------	--------------

AB		SUN PHARM INDUSTRIES	150MG	A072050 001	Apr 17, 1991
-----------	--	-------------------------	--------------	--------------------	--------------

AB			200MG	A072051 001	Apr 17, 1991
-----------	--	--	--------------	--------------------	--------------

AB		WATSON LABS	150MG	A071891 001	Apr 03, 1990
-----------	--	-------------	--------------	--------------------	--------------

AB	!		200MG	A071795 001	Apr 03, 1990
-----------	----------	--	--------------	--------------------	--------------

SUMATRIPTAN

SPRAY; NASAL

IMITREX

AB	+ !	GLAXOSMITHKLINE	5MG/SPRAY	N020626 001	Aug 26, 1997
-----------	------------	-----------------	------------------	--------------------	--------------

AB	+ !		20MG/SPRAY	N020626 003	Aug 26, 1997
-----------	------------	--	-------------------	--------------------	--------------

SUMATRIPTAN

AB		ADAPTIS	20MG/SPRAY	A208967 001	Feb 17, 2021
-----------	--	---------	-------------------	--------------------	--------------

AB		CIPLA	20MG/SPRAY	A214209 001	Feb 22, 2021
-----------	--	-------	-------------------	--------------------	--------------

AB		LANNETT CO INC	5MG/SPRAY	A204841 001	Feb 19, 2016
-----------	--	----------------	------------------	--------------------	--------------

AB			20MG/SPRAY	A204841 002	Feb 19, 2016
-----------	--	--	-------------------	--------------------	--------------

AB		PADAGIS ISRAEL	5MG/SPRAY	A213465 002	Sep 21, 2020
-----------	--	----------------	------------------	--------------------	--------------

AB			20MG/SPRAY	A213465 001	Sep 21, 2020
-----------	--	--	-------------------	--------------------	--------------

TOSYMRA

+ !	TONIX MEDS	10MG/SPRAY	N210884 001	Jan 25, 2019	
------------	------------	-------------------	--------------------	--------------	--

PRESCRIPTION DRUG PRODUCT LIST

SUMATRIPTAN SUCCINATE

INJECTABLE; SUBCUTANEOUS

IMITREX STATDOSE

AB	+	GLAXOSMITHKLINE	<u>EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)</u>	<u>N020080 003</u>	Dec 23, 1996
-----------	----------	-----------------	--	--------------------	--------------

SUMATRIPTAN SUCCINATE

AB		DR REDDYS	<u>EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)</u>	<u>A090495 001</u>	Jan 29, 2014
-----------	--	-----------	--	--------------------	--------------

AP		CAPLIN	<u>EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)</u>	<u>A213998 001</u>	Jul 13, 2021
-----------	--	--------	--	--------------------	--------------

AP	!	EUGIA PHARMA	<u>EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)</u>	<u>A202758 001</u>	Apr 23, 2013
-----------	----------	--------------	--	--------------------	--------------

AP		FRESENIUS KABI USA	<u>EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)</u>	<u>A079242 001</u>	Mar 02, 2009
-----------	--	--------------------	--	--------------------	--------------

AP		HIKMA	<u>EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)</u>	<u>A079123 001</u>	Feb 06, 2009
-----------	--	-------	--	--------------------	--------------

AP			<u>EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)</u>	<u>A200183 001</u>	Sep 16, 2013
-----------	--	--	--	--------------------	--------------

AP		STERISCIENCE	<u>EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)</u>	<u>A215651 001</u>	Nov 21, 2024
-----------	--	--------------	--	--------------------	--------------

AP		SPECLTS			
-----------	--	---------	--	--	--

AP		WOCKHARDT	<u>EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)</u>	<u>A078593 001</u>	Feb 06, 2009
-----------	--	-----------	--	--------------------	--------------

IMITREX STATDOSE

+		GLAXOSMITHKLINE	EQ 4MG BASE/0.5ML (EQ 8MG BASE/ML)	N020080 002	Feb 01, 2006
----------	--	-----------------	------------------------------------	-------------	--------------

POWDER; NASAL

ONZETRA XSAIL

+		CURRAX	EQ 11MG BASE	N206099 001	Jan 27, 2016
----------	--	--------	--------------	-------------	--------------

SOLUTION; SUBCUTANEOUS

ZEMBRACE SYMTOUCH

+		TONIX MEDS	EQ 3MG BASE/0.5ML (EQ 3MG BASE/0.5ML)	N208223 001	Jan 28, 2016
----------	--	------------	---------------------------------------	-------------	--------------

TABLET; ORAL

IMITREX

AB	+	GLAXOSMITHKLINE	<u>EQ 25MG BASE</u>	<u>N020132 002</u>	Jun 01, 1995
-----------	----------	-----------------	---------------------	--------------------	--------------

AB	+		<u>EQ 50MG BASE</u>	<u>N020132 003</u>	Jun 01, 1995
-----------	----------	--	---------------------	--------------------	--------------

AB	+		<u>EQ 100MG BASE</u>	<u>N020132 001</u>	Jun 01, 1995
-----------	----------	--	----------------------	--------------------	--------------

SUMATRIPTAN SUCCINATE

AB		AUROBINDO PHARMA	<u>EQ 25MG BASE</u>	<u>A078327 001</u>	Aug 10, 2009
-----------	--	------------------	---------------------	--------------------	--------------

AB			<u>EQ 50MG BASE</u>	<u>A078327 002</u>	Aug 10, 2009
-----------	--	--	---------------------	--------------------	--------------

AB			<u>EQ 100MG BASE</u>	<u>A078327 003</u>	Aug 10, 2009
-----------	--	--	----------------------	--------------------	--------------

AB		COREPHARMA	<u>EQ 25MG BASE</u>	<u>A200263 001</u>	Jun 19, 2012
-----------	--	------------	---------------------	--------------------	--------------

AB			<u>EQ 50MG BASE</u>	<u>A200263 002</u>	Jun 19, 2012
-----------	--	--	---------------------	--------------------	--------------

AB			<u>EQ 100MG BASE</u>	<u>A200263 003</u>	Jun 19, 2012
-----------	--	--	----------------------	--------------------	--------------

AB		DR REDDYS LABS INC	<u>EQ 25MG BASE</u>	<u>A076847 001</u>	Aug 10, 2009
-----------	--	--------------------	---------------------	--------------------	--------------

AB			<u>EQ 50MG BASE</u>	<u>A076847 002</u>	Aug 10, 2009
-----------	--	--	---------------------	--------------------	--------------

AB			<u>EQ 100MG BASE</u>	<u>A076847 003</u>	Aug 10, 2009
-----------	--	--	----------------------	--------------------	--------------

AB		MYLAN	<u>EQ 25MG BASE</u>	<u>A077744 001</u>	Aug 10, 2009
-----------	--	-------	---------------------	--------------------	--------------

AB			<u>EQ 50MG BASE</u>	<u>A077744 002</u>	Aug 10, 2009
-----------	--	--	---------------------	--------------------	--------------

AB			<u>EQ 100MG BASE</u>	<u>A077744 003</u>	Aug 10, 2009
-----------	--	--	----------------------	--------------------	--------------

AB		ORBION PHARMS	<u>EQ 25MG BASE</u>	<u>A078284 001</u>	Aug 10, 2009
-----------	--	---------------	---------------------	--------------------	--------------

AB			<u>EQ 50MG BASE</u>	<u>A078284 002</u>	Aug 10, 2009
-----------	--	--	---------------------	--------------------	--------------

AB			<u>EQ 100MG BASE</u>	<u>A078284 003</u>	Aug 10, 2009
-----------	--	--	----------------------	--------------------	--------------

AB		SUN PHARM INDS	<u>EQ 25MG BASE</u>	<u>A078295 001</u>	Aug 10, 2009
-----------	--	----------------	---------------------	--------------------	--------------

AB			<u>EQ 50MG BASE</u>	<u>A078295 002</u>	Aug 10, 2009
-----------	--	--	---------------------	--------------------	--------------

AB			<u>EQ 100MG BASE</u>	<u>A078295 003</u>	Aug 10, 2009
-----------	--	--	----------------------	--------------------	--------------

AB		VKT PHARMA	<u>EQ 25MG BASE</u>	<u>A219036 001</u>	Dec 31, 2024
-----------	--	------------	---------------------	--------------------	--------------

AB			<u>EQ 50MG BASE</u>	<u>A219036 002</u>	Dec 31, 2024
-----------	--	--	---------------------	--------------------	--------------

AB			<u>EQ 100MG BASE</u>	<u>A219036 003</u>	Dec 31, 2024
-----------	--	--	----------------------	--------------------	--------------

AB		WATSON LABS	<u>EQ 25MG BASE</u>	<u>A076933 001</u>	Aug 10, 2009
-----------	--	-------------	---------------------	--------------------	--------------

AB			<u>EQ 50MG BASE</u>	<u>A076933 002</u>	Aug 10, 2009
-----------	--	--	---------------------	--------------------	--------------

AB			<u>EQ 100MG BASE</u>	<u>A076933 003</u>	Aug 10, 2009
-----------	--	--	----------------------	--------------------	--------------

SUNITINIB MALATE

CAPSULE; ORAL

SUNITINIB MALATE

AB		DR REDDYS	<u>EQ 12.5MG BASE</u>	<u>A215843 001</u>	Apr 11, 2022
-----------	--	-----------	-----------------------	--------------------	--------------

AB			<u>EQ 25MG BASE</u>	<u>A215843 002</u>	Apr 11, 2022
-----------	--	--	---------------------	--------------------	--------------

AB			<u>EQ 37.5MG BASE</u>	<u>A215843 003</u>	Apr 11, 2022
-----------	--	--	-----------------------	--------------------	--------------

AB			<u>EQ 50MG BASE</u>	<u>A215843 004</u>	Apr 11, 2022
-----------	--	--	---------------------	--------------------	--------------

AB		EUGIA PHARMA	<u>EQ 12.5MG BASE</u>	<u>A218615 001</u>	Mar 14, 2024
-----------	--	--------------	-----------------------	--------------------	--------------

AB			<u>EQ 25MG BASE</u>	<u>A218615 002</u>	Mar 14, 2024
-----------	--	--	---------------------	--------------------	--------------

AB			<u>EQ 37.5MG BASE</u>	<u>A218615 003</u>	Mar 14, 2024
-----------	--	--	-----------------------	--------------------	--------------

AB			<u>EQ 50MG BASE</u>	<u>A218615 004</u>	Mar 14, 2024
-----------	--	--	---------------------	--------------------	--------------

AB		MYLAN	<u>EQ 12.5MG BASE</u>	<u>A201275 001</u>	Dec 06, 2021
-----------	--	-------	-----------------------	--------------------	--------------

AB			<u>EQ 25MG BASE</u>	<u>A201275 002</u>	Dec 06, 2021
-----------	--	--	---------------------	--------------------	--------------

AB			<u>EQ 37.5MG BASE</u>	<u>A201275 003</u>	Dec 06, 2021
-----------	--	--	-----------------------	--------------------	--------------

AB			<u>EQ 50MG BASE</u>	<u>A201275 004</u>	Dec 06, 2021
-----------	--	--	---------------------	--------------------	--------------

AB		NATCO PHARMA	<u>EQ 12.5MG BASE</u>	<u>A218024 001</u>	Oct 24, 2023
-----------	--	--------------	-----------------------	--------------------	--------------

AB			<u>EQ 25MG BASE</u>	<u>A218024 002</u>	Oct 24, 2023
-----------	--	--	---------------------	--------------------	--------------

AB			<u>EQ 37.5MG BASE</u>	<u>A218024 003</u>	Oct 24, 2023
-----------	--	--	-----------------------	--------------------	--------------

AB			<u>EQ 50MG BASE</u>	<u>A218024 004</u>	Oct 24, 2023
-----------	--	--	---------------------	--------------------	--------------

AB		SUN PHARM	<u>EQ 12.5MG BASE</u>	<u>A213914 001</u>	Aug 16, 2021
-----------	--	-----------	-----------------------	--------------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

SUNITINIB MALATE

CAPSULE; ORAL

SUNITINIB MALATE

<u>AB</u>		<u>EQ 25MG BASE</u>	<u>A213914 002</u>	Aug 16, 2021
<u>AB</u>		<u>EQ 37.5MG BASE</u>	<u>A213914 003</u>	Aug 16, 2021
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A213914 004</u>	Aug 16, 2021
<u>AB</u>	TEVA PHARMS USA	<u>EQ 12.5MG BASE</u>	<u>A213803 001</u>	Nov 30, 2021
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>A213803 002</u>	Nov 30, 2021
<u>AB</u>		<u>EQ 37.5MG BASE</u>	<u>A213803 003</u>	Nov 30, 2021
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A213803 004</u>	Nov 30, 2021
<u>AB</u>	WANBANG BIOPHARMS	<u>EQ 12.5MG BASE</u>	<u>A218012 001</u>	Aug 21, 2023
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>A218012 002</u>	Aug 21, 2023
<u>AB</u>		<u>EQ 37.5MG BASE</u>	<u>A218012 003</u>	Aug 21, 2023
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A218012 004</u>	Aug 21, 2023

SUTENT

<u>AB</u>	+	CPPI CV	<u>EQ 12.5MG BASE</u>	<u>N021938 001</u>	Jan 26, 2006
<u>AB</u>	+		<u>EQ 25MG BASE</u>	<u>N021938 002</u>	Jan 26, 2006
<u>AB</u>	+		<u>EQ 37.5MG BASE</u>	<u>N021938 004</u>	Mar 31, 2009
<u>AB</u>	+	!	<u>EQ 50MG BASE</u>	<u>N021938 003</u>	Jan 26, 2006

SUVOREXANT

TABLET; ORAL

BELSOMRA

+	MERCK SHARP DOHME	5MG	N204569 001	Aug 13, 2014
+		10MG	N204569 002	Aug 13, 2014
+		15MG	N204569 003	Aug 13, 2014
+	!	20MG	N204569 004	Aug 13, 2014

TACROLIMUS

CAPSULE; ORAL

PROGRAF

<u>AB</u>	+	ASTELLAS	<u>EQ 0.5MG BASE</u>	<u>N050708 003</u>	Aug 24, 1998
<u>AB</u>	+		<u>EQ 1MG BASE</u>	<u>N050708 001</u>	Apr 08, 1994
<u>AB</u>	+	!	<u>EQ 5MG BASE</u>	<u>N050708 002</u>	Apr 08, 1994

TACROLIMUS

<u>AB</u>		ALKEM LABS LTD	<u>EQ 0.5MG BASE</u>	<u>A203740 001</u>	Nov 12, 2020
<u>AB</u>			<u>EQ 1MG BASE</u>	<u>A203740 002</u>	Nov 12, 2020
<u>AB</u>			<u>EQ 5MG BASE</u>	<u>A203740 003</u>	Nov 12, 2020
<u>AB</u>		BELCHER	<u>EQ 0.5MG BASE</u>	<u>A206651 001</u>	Nov 30, 2017
<u>AB</u>			<u>EQ 1MG BASE</u>	<u>A206651 002</u>	Nov 30, 2017
<u>AB</u>			<u>EQ 5MG BASE</u>	<u>A206651 003</u>	Nov 30, 2017
<u>AB</u>		BIOCON PHARMA	<u>EQ 0.5MG BASE</u>	<u>A212297 001</u>	Nov 10, 2020
<u>AB</u>			<u>EQ 1MG BASE</u>	<u>A212297 002</u>	Nov 10, 2020
<u>AB</u>			<u>EQ 5MG BASE</u>	<u>A212297 003</u>	Nov 10, 2020
<u>AB</u>		CONCORD BIOTECH LTD	<u>EQ 0.5MG BASE</u>	<u>A213112 001</u>	Nov 10, 2020
<u>AB</u>			<u>EQ 1MG BASE</u>	<u>A213112 002</u>	Nov 10, 2020
<u>AB</u>			<u>EQ 5MG BASE</u>	<u>A213112 003</u>	Nov 10, 2020
<u>AB</u>		DR REDDYS LABS LTD	<u>EQ 0.5MG BASE</u>	<u>A090509 001</u>	May 12, 2010
<u>AB</u>			<u>EQ 1MG BASE</u>	<u>A090509 002</u>	May 12, 2010
<u>AB</u>			<u>EQ 5MG BASE</u>	<u>A090509 003</u>	May 12, 2010
<u>AB</u>		GLENMARK PHARMS LTD	<u>EQ 0.5MG BASE</u>	<u>A206662 001</u>	Nov 10, 2020
<u>AB</u>			<u>EQ 1MG BASE</u>	<u>A206662 002</u>	Nov 10, 2020
<u>AB</u>			<u>EQ 5MG BASE</u>	<u>A206662 003</u>	Nov 10, 2020
<u>AB</u>		HANGZHOU ZHONGMEI	<u>EQ 0.5MG BASE</u>	<u>A210929 001</u>	Apr 17, 2023
<u>AB</u>			<u>EQ 1MG BASE</u>	<u>A210929 002</u>	Apr 17, 2023
<u>AB</u>			<u>EQ 5MG BASE</u>	<u>A210929 003</u>	Apr 17, 2023
<u>AB</u>		MYLAN	<u>EQ 0.5MG BASE</u>	<u>A090596 001</u>	Sep 17, 2010
<u>AB</u>			<u>EQ 1MG BASE</u>	<u>A090596 002</u>	Sep 17, 2010
<u>AB</u>			<u>EQ 5MG BASE</u>	<u>A090596 003</u>	Sep 17, 2010
<u>AB</u>		PANACEA	<u>EQ 0.5MG BASE</u>	<u>A090802 001</u>	Sep 28, 2012
<u>AB</u>			<u>EQ 1MG BASE</u>	<u>A090802 002</u>	Sep 28, 2012
<u>AB</u>			<u>EQ 5MG BASE</u>	<u>A090802 003</u>	Sep 28, 2012
<u>AB</u>		SANDOZ	<u>EQ 0.5MG BASE</u>	<u>A065461 001</u>	Aug 10, 2009
<u>AB</u>			<u>EQ 1MG BASE</u>	<u>A065461 002</u>	Aug 10, 2009
<u>AB</u>			<u>EQ 5MG BASE</u>	<u>A065461 003</u>	Aug 10, 2009
<u>AB</u>		STRIDES PHARMA	<u>EQ 0.5MG BASE</u>	<u>A090687 001</u>	Jul 22, 2014
<u>AB</u>			<u>EQ 1MG BASE</u>	<u>A090687 002</u>	Jul 22, 2014
<u>AB</u>			<u>EQ 5MG BASE</u>	<u>A090687 003</u>	Jul 22, 2014
BX		ACCORD HLTHCARE	EQ 0.5MG BASE	A091195 001	Aug 31, 2011
BX			EQ 1MG BASE	A091195 002	Aug 31, 2011
BX			EQ 5MG BASE	A091195 003	Aug 31, 2011

PRESCRIPTION DRUG PRODUCT LIST

TACROLIMUS

CAPSULE, EXTENDED RELEASE;ORAL

ASTAGRAF XL

<u>AB</u>	+	ASTELLAS	<u>EQ 0.5MG BASE</u>	<u>N204096 001</u>	Jul 19, 2013
<u>AB</u>	+		<u>EQ 1MG BASE</u>	<u>N204096 002</u>	Jul 19, 2013
<u>AB</u>	+	!	<u>EQ 5MG BASE</u>	<u>N204096 003</u>	Jul 19, 2013

TACROLIMUS

<u>AB</u>		CHENGDU	<u>EQ 0.5MG BASE</u>	<u>A215012 001</u>	Jan 25, 2024
<u>AB</u>			<u>EQ 1MG BASE</u>	<u>A215012 002</u>	Jan 25, 2024
<u>AB</u>			<u>EQ 5MG BASE</u>	<u>A215012 003</u>	Jan 25, 2024

FOR SUSPENSION;ORAL

PROGRAF

	+	ASTELLAS	EQ 0.2MG BASE/PACKET	N210115 001	May 24, 2018
	+	!	EQ 1MG BASE/PACKET	N210115 002	May 24, 2018

INJECTABLE; INJECTION

PROGRAF

	+	!	ASTELLAS	EQ 5MG BASE/ML	N050709 001	Apr 08, 1994
--	---	---	----------	----------------	-------------	--------------

OINTMENT; TOPICAL

PROTOPIC

<u>AB</u>	+	!	LEO PHARMA AS	<u>0.03%</u>	<u>N050777 001</u>	Dec 08, 2000
<u>AB</u>	+	!		<u>0.1%</u>	<u>N050777 002</u>	Dec 08, 2000

TACROLIMUS

<u>AB</u>		ACCORD HLTHCARE	<u>0.03%</u>	<u>A211688 001</u>	Jan 31, 2019
<u>AB</u>			<u>0.1%</u>	<u>A211688 002</u>	Jan 31, 2019
<u>AB</u>		ENCUBE	<u>0.1%</u>	<u>A212387 001</u>	Oct 10, 2023
<u>AB</u>		FOUGERA PHARMS INC	<u>0.03%</u>	<u>A200744 001</u>	Sep 09, 2014
<u>AB</u>			<u>0.1%</u>	<u>A200744 002</u>	Sep 09, 2014
<u>AB</u>		GLENMARK PHARMS LTD	<u>0.03%</u>	<u>A210393 002</u>	Aug 16, 2023
<u>AB</u>			<u>0.1%</u>	<u>A210393 001</u>	Apr 16, 2018

TABLET, EXTENDED RELEASE;ORAL

ENVARUSUS XR

	+	VELOXIS PHARMS INC	EQ 0.75MG BASE	N206406 001	Jul 10, 2015
	+		EQ 1MG BASE	N206406 002	Jul 10, 2015
	+	!	EQ 4MG BASE	N206406 003	Jul 10, 2015

TADALAFIL

SUSPENSION; ORAL

TADLIQ

	+	!	CMP DEV LLC	20MG/5ML	N214522 001	Jun 17, 2022
--	---	---	-------------	----------	-------------	--------------

TABLET; ORAL

CIALIS

<u>AB1</u>	+	LILLY	<u>5MG</u>	<u>N021368 001</u>	Nov 21, 2003
<u>AB1</u>	+		<u>10MG</u>	<u>N021368 002</u>	Nov 21, 2003
<u>AB1</u>	+	!	<u>20MG</u>	<u>N021368 003</u>	Nov 21, 2003

TADALAFIL

<u>AB1</u>		ACCORD HLTHCARE	<u>2.5MG</u>	<u>A209167 001</u>	Mar 26, 2019
<u>AB1</u>			<u>5MG</u>	<u>A209167 002</u>	Mar 26, 2019
<u>AB1</u>			<u>10MG</u>	<u>A209167 003</u>	Oct 23, 2019
<u>AB1</u>			<u>20MG</u>	<u>A209167 004</u>	Mar 26, 2019
<u>AB1</u>		AJANTA PHARMA LTD	<u>2.5MG</u>	<u>A209654 001</u>	Mar 26, 2019
<u>AB1</u>			<u>5MG</u>	<u>A209654 002</u>	Mar 26, 2019
<u>AB1</u>			<u>10MG</u>	<u>A209654 003</u>	Mar 26, 2019
<u>AB1</u>			<u>20MG</u>	<u>A209654 004</u>	Mar 26, 2019
<u>AB1</u>		ALEMBIC	<u>2.5MG</u>	<u>A204809 001</u>	Mar 26, 2019
<u>AB1</u>			<u>5MG</u>	<u>A204809 002</u>	Mar 26, 2019
<u>AB1</u>			<u>10MG</u>	<u>A204809 003</u>	Mar 26, 2019
<u>AB1</u>			<u>20MG</u>	<u>A204809 004</u>	Mar 26, 2019
<u>AB1</u>		AMNEAL PHARMS CO	<u>2.5MG</u>	<u>A209744 001</u>	Mar 26, 2019
<u>AB1</u>			<u>5MG</u>	<u>A209744 002</u>	Mar 26, 2019
<u>AB1</u>			<u>10MG</u>	<u>A209744 003</u>	Mar 26, 2019
<u>AB1</u>			<u>20MG</u>	<u>A209744 004</u>	Mar 26, 2019
<u>AB1</u>		AUROBINDO PHARMA LTD	<u>2.5MG</u>	<u>A206285 001</u>	Mar 26, 2019
<u>AB1</u>			<u>5MG</u>	<u>A206285 002</u>	Mar 26, 2019
<u>AB1</u>			<u>10MG</u>	<u>A206285 003</u>	Mar 26, 2019
<u>AB1</u>			<u>20MG</u>	<u>A206285 004</u>	Mar 26, 2019
<u>AB1</u>		CIPLA	<u>2.5MG</u>	<u>A209539 001</u>	Mar 26, 2019
<u>AB1</u>			<u>5MG</u>	<u>A209539 002</u>	Mar 26, 2019
<u>AB1</u>			<u>10MG</u>	<u>A209539 003</u>	Mar 26, 2019
<u>AB1</u>			<u>20MG</u>	<u>A209539 004</u>	Mar 26, 2019
<u>AB1</u>		DR REDDYS	<u>2.5MG</u>	<u>A210069 001</u>	Mar 26, 2019
<u>AB1</u>			<u>5MG</u>	<u>A210069 002</u>	Mar 26, 2019
<u>AB1</u>			<u>10MG</u>	<u>A210069 003</u>	Mar 26, 2019

PRESCRIPTION DRUG PRODUCT LIST

TADALAFIL

TABLET; ORAL

TADALAFIL

<u>AB1</u>		<u>20MG</u>	<u>A210069 004</u>	Mar 26, 2019
<u>AB1</u>	FOURRTS LABS	<u>2.5MG</u>	<u>A217606 001</u>	Aug 13, 2024
<u>AB1</u>		<u>5MG</u>	<u>A217606 002</u>	Aug 13, 2024
<u>AB1</u>		<u>10MG</u>	<u>A217606 003</u>	Aug 13, 2024
<u>AB1</u>		<u>20MG</u>	<u>A217606 004</u>	Aug 13, 2024
<u>AB1</u>	HANGZHOU BINJIANG	<u>2.5MG</u>	<u>A208824 001</u>	Oct 27, 2020
<u>AB1</u>		<u>5MG</u>	<u>A208824 002</u>	Oct 27, 2020
<u>AB1</u>		<u>10MG</u>	<u>A208824 003</u>	Oct 27, 2020
<u>AB1</u>		<u>20MG</u>	<u>A208824 004</u>	Oct 27, 2020
<u>AB1</u>	HETERO LABS LTD III	<u>2.5MG</u>	<u>A209908 001</u>	Mar 26, 2019
<u>AB1</u>		<u>5MG</u>	<u>A209908 002</u>	Mar 26, 2019
<u>AB1</u>		<u>10MG</u>	<u>A209908 003</u>	Mar 26, 2019
<u>AB1</u>		<u>20MG</u>	<u>A209908 004</u>	Mar 26, 2019
<u>AB1</u>	LUPIN LTD	<u>2.5MG</u>	<u>A210567 001</u>	Mar 26, 2019
<u>AB1</u>		<u>5MG</u>	<u>A210567 002</u>	Mar 26, 2019
<u>AB1</u>		<u>10MG</u>	<u>A210567 003</u>	Mar 26, 2019
<u>AB1</u>		<u>20MG</u>	<u>A210567 004</u>	Mar 26, 2019
<u>AB1</u>	MACLEODS PHARMS LTD	<u>2.5MG</u>	<u>A207244 001</u>	Oct 07, 2019
<u>AB1</u>		<u>5MG</u>	<u>A207244 002</u>	Oct 07, 2019
<u>AB1</u>		<u>10MG</u>	<u>A207244 003</u>	Oct 07, 2019
<u>AB1</u>		<u>20MG</u>	<u>A207244 004</u>	Oct 07, 2019
<u>AB1</u>	NOVITIUM PHARMA	<u>5MG</u>	<u>A215949 001</u>	Mar 03, 2023
<u>AB1</u>		<u>10MG</u>	<u>A215949 002</u>	Mar 03, 2023
<u>AB1</u>		<u>20MG</u>	<u>A215949 003</u>	Mar 03, 2023
<u>AB1</u>	PRINSTON INC	<u>2.5MG</u>	<u>A210609 001</u>	Aug 11, 2022
<u>AB1</u>		<u>5MG</u>	<u>A210609 002</u>	Aug 11, 2022
<u>AB1</u>		<u>10MG</u>	<u>A210609 003</u>	Aug 11, 2022
<u>AB1</u>		<u>20MG</u>	<u>A210609 004</u>	Aug 11, 2022
<u>AB1</u>	QILU PHARM HAINAN	<u>2.5MG</u>	<u>A210420 001</u>	Mar 26, 2019
<u>AB1</u>		<u>5MG</u>	<u>A210420 002</u>	Mar 26, 2019
<u>AB1</u>		<u>10MG</u>	<u>A210420 003</u>	Mar 26, 2019
<u>AB1</u>		<u>20MG</u>	<u>A210420 004</u>	Mar 26, 2019
<u>AB1</u>	SHANDONG	<u>2.5MG</u>	<u>A216279 002</u>	Dec 04, 2024
<u>AB1</u>		<u>5MG</u>	<u>A216279 003</u>	Dec 04, 2024
<u>AB1</u>		<u>10MG</u>	<u>A216279 004</u>	Dec 04, 2024
<u>AB1</u>		<u>20MG</u>	<u>A216279 001</u>	Mar 04, 2024
<u>AB1</u>	SUN PHARM	<u>2.5MG</u>	<u>A208934 001</u>	Mar 26, 2019
<u>AB1</u>		<u>5MG</u>	<u>A208934 002</u>	Mar 26, 2019
<u>AB1</u>		<u>10MG</u>	<u>A208934 003</u>	Mar 26, 2019
<u>AB1</u>		<u>20MG</u>	<u>A208934 004</u>	Mar 26, 2019
<u>AB1</u>	SUNSHINE	<u>2.5MG</u>	<u>A211335 001</u>	Mar 26, 2019
<u>AB1</u>		<u>5MG</u>	<u>A211335 002</u>	Mar 26, 2019
<u>AB1</u>		<u>10MG</u>	<u>A211335 003</u>	Mar 26, 2019
<u>AB1</u>		<u>20MG</u>	<u>A211335 004</u>	Mar 26, 2019
<u>AB1</u>	TEVA PHARMS USA	<u>2.5MG</u>	<u>A090141 001</u>	May 22, 2018
<u>AB1</u>		<u>5MG</u>	<u>A090141 002</u>	May 22, 2018
<u>AB1</u>		<u>10MG</u>	<u>A090141 003</u>	May 22, 2018
<u>AB1</u>		<u>20MG</u>	<u>A090141 004</u>	May 22, 2018
<u>AB1</u>	TORRENT	<u>2.5MG</u>	<u>A211839 001</u>	Mar 26, 2019
<u>AB1</u>		<u>5MG</u>	<u>A211839 002</u>	Mar 26, 2019
<u>AB1</u>		<u>10MG</u>	<u>A211839 003</u>	Mar 26, 2019
<u>AB1</u>		<u>20MG</u>	<u>A211839 004</u>	Mar 26, 2019
<u>AB1</u>	UNICHEM	<u>2.5MG</u>	<u>A209250 001</u>	Mar 26, 2019
<u>AB1</u>		<u>5MG</u>	<u>A209250 002</u>	Mar 26, 2019
<u>AB1</u>		<u>10MG</u>	<u>A209250 003</u>	Mar 26, 2019
<u>AB1</u>		<u>20MG</u>	<u>A209250 004</u>	Mar 26, 2019
<u>AB1</u>	VKT PHARMA	<u>2.5MG</u>	<u>A215556 001</u>	Nov 04, 2021
<u>AB1</u>		<u>5MG</u>	<u>A215556 002</u>	Nov 04, 2021
<u>AB1</u>		<u>10MG</u>	<u>A215556 003</u>	Nov 04, 2021
<u>AB1</u>		<u>20MG</u>	<u>A215556 004</u>	Nov 04, 2021
<u>AB1</u>	WATSON LABS INC	<u>2.5MG</u>	<u>A205885 001</u>	Mar 29, 2019
<u>AB1</u>		<u>5MG</u>	<u>A205885 002</u>	Mar 29, 2019
<u>AB1</u>		<u>10MG</u>	<u>A205885 003</u>	Mar 29, 2019
<u>AB1</u>		<u>20MG</u>	<u>A205885 004</u>	Mar 29, 2019
<u>AB1</u>	ZYDUS PHARMS	<u>2.5MG</u>	<u>A206693 001</u>	Mar 26, 2019
<u>AB1</u>		<u>5MG</u>	<u>A206693 002</u>	Mar 26, 2019
<u>AB1</u>		<u>10MG</u>	<u>A206693 003</u>	Mar 26, 2019
<u>AB1</u>		<u>20MG</u>	<u>A206693 004</u>	Mar 26, 2019

PRESCRIPTION DRUG PRODUCT LIST

TADALAFIL

TABLET; ORAL

ADCIRCA

AB2	+ !	ELI LILLY CO	20MG	N022332	001	May 22, 2009
------------	------------	--------------	-------------	----------------	------------	--------------

ALYO

AB2		TEVA PHARMS INC	20MG	A216932	001	Oct 12, 2022
------------	--	-----------------	-------------	----------------	------------	--------------

AB2		TEVA PHARMS USA	20MG	A209942	001	Feb 05, 2019
------------	--	-----------------	-------------	----------------	------------	--------------

TADALAFIL

AB2		AJANTA PHARMA LTD	20MG	A210392	001	Feb 05, 2019
------------	--	-------------------	-------------	----------------	------------	--------------

AB2		AUROBINDO PHARMA LTD	20MG	A206286	001	Feb 05, 2019
------------	--	----------------------	-------------	----------------	------------	--------------

AB2		CHARTWELL RX	20MG	A210572	001	Feb 05, 2019
------------	--	--------------	-------------	----------------	------------	--------------

AB2		CIPLA	20MG	A210255	001	Feb 05, 2019
------------	--	-------	-------------	----------------	------------	--------------

AB2		DR REDDYS	20MG	A210145	001	Feb 05, 2019
------------	--	-----------	-------------	----------------	------------	--------------

AB2		HETERO LABS LTD III	20MG	A209907	001	Feb 05, 2019
------------	--	---------------------	-------------	----------------	------------	--------------

AB2		MACLEODS PHARMS LTD	20MG	A207290	001	Oct 16, 2019
------------	--	---------------------	-------------	----------------	------------	--------------

AB2		PRINSTON INC	20MG	A210608	001	Aug 11, 2022
------------	--	--------------	-------------	----------------	------------	--------------

AB2		SUNSHINE	20MG	A213496	001	Nov 23, 2020
------------	--	----------	-------------	----------------	------------	--------------

AB2		TORRENT	20MG	A212062	001	Mar 26, 2019
------------	--	---------	-------------	----------------	------------	--------------

BX		UMEDICA	2.5MG	A211298	001	Oct 23, 2020
----	--	---------	-------	---------	-----	--------------

BX			5MG	A211298	002	Oct 23, 2020
----	--	--	-----	---------	-----	--------------

BX			10MG	A211298	003	Oct 23, 2020
----	--	--	------	---------	-----	--------------

BX			20MG	A211298	004	Oct 23, 2020
----	--	--	------	---------	-----	--------------

TAFAMIDIS

CAPSULE; ORAL

VYNDAMAX

+ !		FOLDRX PHARMS	61MG	N212161	001	May 03, 2019
------------	--	---------------	------	---------	-----	--------------

TAFAMIDIS MEGLUMINE

CAPSULE; ORAL

VYNDAQEL

+ !		FOLDRX PHARMS	20MG	N211996	001	May 03, 2019
------------	--	---------------	------	---------	-----	--------------

TAFENOQUINE SUCCINATE

TABLET; ORAL

ARAKODA

+ !		60 DEGREES PHARMS	EQ 100MG BASE	N210607	001	Aug 08, 2018
------------	--	-------------------	---------------	---------	-----	--------------

KRINTAFEL

+ !		GLAXOSMITHKLINE	EQ 150MG BASE	N210795	001	Jul 20, 2018
------------	--	-----------------	---------------	---------	-----	--------------

TAFLUPROST

SOLUTION/DROPS; OPHTHALMIC

TAFLUPROST

AT		INGENUS PHARMS LLC	0.0015%	A218002	001	Apr 05, 2024
-----------	--	--------------------	----------------	----------------	------------	--------------

AT		MICRO LABS	0.0015%	A209051	001	Aug 19, 2019
-----------	--	------------	----------------	----------------	------------	--------------

AT		SANDOZ	0.0015%	A209040	001	Jan 28, 2022
-----------	--	--------	----------------	----------------	------------	--------------

ZIOPTAN

AT	+ !	THEA PHARMA	0.0015%	N202514	001	Feb 10, 2012
-----------	------------	-------------	----------------	----------------	------------	--------------

TALAZOPARIB TOSYLATE

CAPSULE; ORAL

TALZENNA

+		PFIZER	EQ 0.1MG BASE	N211651	005	Jun 20, 2023
----------	--	--------	---------------	---------	-----	--------------

+			EQ 0.1MG BASE	N217439	001	Mar 07, 2024
----------	--	--	---------------	---------	-----	--------------

+			EQ 0.25MG BASE	N211651	001	Oct 16, 2018
----------	--	--	----------------	---------	-----	--------------

+			EQ 0.25MG BASE	N217439	002	Mar 07, 2024
----------	--	--	----------------	---------	-----	--------------

+			EQ 0.35MG BASE	N211651	006	Jun 20, 2023
----------	--	--	----------------	---------	-----	--------------

+			EQ 0.35MG BASE	N217439	003	Mar 07, 2024
----------	--	--	----------------	---------	-----	--------------

+			EQ 0.5MG BASE	N211651	003	Sep 20, 2021
----------	--	--	---------------	---------	-----	--------------

+			EQ 0.5MG BASE	N217439	004	Mar 07, 2024
----------	--	--	---------------	---------	-----	--------------

+			EQ 0.75MG BASE	N211651	004	Sep 20, 2021
----------	--	--	----------------	---------	-----	--------------

+			EQ 0.75MG BASE	N217439	005	Mar 07, 2024
----------	--	--	----------------	---------	-----	--------------

+ !			EQ 1MG BASE	N211651	002	Oct 16, 2018
------------	--	--	-------------	---------	-----	--------------

+ !			EQ 1MG BASE	N217439	006	Mar 07, 2024
------------	--	--	-------------	---------	-----	--------------

TALC

AEROSOL; INTRAPLEURAL

SCLEROSOL

+ !		SCIARRA LABS	4GM/SPRAY	N020587	001	Dec 24, 1997
------------	--	--------------	-----------	---------	-----	--------------

POWDER; INTRAPLEURAL

STERITALC

+		NOVATECH SA	2GM/VIAL	N205555	001	May 01, 2017
----------	--	-------------	----------	---------	-----	--------------

+			3GM/VIAL	N205555	002	May 01, 2017
----------	--	--	----------	---------	-----	--------------

+ !			4GM/VIAL	N205555	003	May 01, 2017
------------	--	--	----------	---------	-----	--------------

PRESCRIPTION DRUG PRODUCT LIST

TALC

POWDER; INTRAPLEURAL

TALC

+! SCIARRA LABS 5GM/BOT N021388 001 Dec 15, 2003

TAMOXIFEN CITRATE

SOLUTION; ORAL

SOLTAMOX

+! MAYNE PHARMA EQ 20MG BASE/10ML N021807 001 Oct 29, 2005
COMMCL

TABLET; ORAL

TAMOXIFEN CITRATE

AB	ACTAVIS LABS FL INC	EQ 10MG BASE	A070929 001	Feb 20, 2003
AB		EQ 20MG BASE	A070929 002	Feb 20, 2003
AB	APOTEX	EQ 10MG BASE	A090878 001	Sep 23, 2011
AB		EQ 20MG BASE	A090878 002	Sep 23, 2011
AB	DR REDDYS LABS SA	EQ 10MG BASE	A075797 001	Feb 20, 2003
AB	!	EQ 20MG BASE	A075797 002	Feb 20, 2003
AB	EUGIA PHARMA	EQ 10MG BASE	A213358 001	Aug 14, 2020
AB		EQ 20MG BASE	A213358 002	Aug 14, 2020
AB	MYLAN	EQ 10MG BASE	A074732 002	Feb 20, 2003
AB		EQ 20MG BASE	A074732 001	Feb 20, 2003
AB	ZYDUS PHARMS	EQ 10MG BASE	A206694 001	Oct 27, 2017
AB		EQ 20MG BASE	A206694 002	Oct 27, 2017

TAMSULOSIN HYDROCHLORIDE

CAPSULE; ORAL

FLOMAX**AB** +! SANOFI **0.4MG** **N020579 001** Apr 15, 1997TAMSULOSIN HYDROCHLORIDE

AB	ALKEM LABS LTD	0.4MG	A207405 001	Aug 11, 2017
AB	AUROBINDO PHARMA LTD	0.4MG	A202433 001	Apr 30, 2013
AB	CHARTWELL RX	0.4MG	A211885 001	Oct 17, 2019
AB	IMPAX LABS	0.4MG	A090377 001	Mar 02, 2010
AB	MACLEODS PHARMS LTD	0.4MG	A204645 001	Jan 20, 2017
AB	SANDOZ	0.4MG	A078015 001	Apr 27, 2010
AB	SCIEGEN PHARMS INC	0.4MG	A078938 001	Apr 27, 2010
AB	SYNTHON BV	0.4MG	A078801 001	Apr 27, 2010
AB	TEVA PHARMS	0.4MG	A077630 001	Apr 27, 2010
AB	ZYDUS PHARMS USA INC	0.4MG	A078225 001	Apr 27, 2010

TAPENTADOL HYDROCHLORIDE

TABLET; ORAL

NUCYNTA

+ COLLEGIUM PHARM INC EQ 50MG BASE N022304 001 Nov 20, 2008
+ EQ 75MG BASE N022304 002 Nov 20, 2008
+! EQ 100MG BASE N022304 003 Nov 20, 2008

TABLET, EXTENDED RELEASE; ORAL

NUCYNTA ER

+ COLLEGIUM PHARM INC EQ 50MG BASE N200533 001 Aug 25, 2011
+ EQ 100MG BASE N200533 002 Aug 25, 2011
+ EQ 150MG BASE N200533 003 Aug 25, 2011
+ EQ 200MG BASE N200533 004 Aug 25, 2011
+! EQ 250MG BASE N200533 005 Aug 25, 2011TAPINAROF

CREAM; TOPICAL

VTAMA

+! DERMAVANT SCI 1% N215272 001 May 23, 2022

TASIMELTEON

CAPSULE; ORAL

HETLIOZ**AB** +! VANDA PHARMS INC **20MG** **N205677 001** Jan 31, 2014TASIMELTEON

AB	APOTEX	20MG	A211607 001	Dec 20, 2022
AB	MSN	20MG	A211654 001	Jan 12, 2023
AB	TEVA PHARMS USA INC	20MG	A211601 001	Dec 12, 2022

SUSPENSION; ORAL

HETLIOZ LQ

+! VANDA PHARMS INC 4MG/ML N214517 001 Dec 01, 2020

PRESCRIPTION DRUG PRODUCT LIST

TAVABOROLE

SOLUTION; TOPICAL

TAVABOROLE

AB	ALEMBIC	5%	A212188 001	Oct 21, 2020
AB	AMNEAL	5%	A212256 001	Nov 25, 2020
AB	CHARTWELL RX	5%	A211963 001	Feb 03, 2021
AB	CIPLA	5%	A212224 001	Feb 09, 2021
AB	! ENCUBE	5%	A211297 001	Oct 13, 2020
AB	LUPIN LTD	5%	A212168 001	Feb 08, 2021
AB	PADAGIS US	5%	A211848 001	Oct 13, 2020
AB	TARO	5%	A212215 001	May 07, 2021
AB	ZYDUS LIFESCIENCES	5%	A212294 001	Apr 10, 2023

TAZAROTENE

AEROSOL, FOAM; TOPICAL

FABIOR

+! MAYNE PHARMA 0.1%

N202428 001 May 11, 2012

CREAM; TOPICAL

AVAGE

AB	+! ALLERGAN	0.1%	N021184 003	Sep 30, 2002
-----------	-------------	-------------	--------------------	--------------

TAZAROTENE

AB	COSETTE	0.1%	A208662 001	Dec 22, 2017
AB	PADAGIS ISRAEL	0.05%	A217075 001	Jul 15, 2024
AB	TARO	0.1%	A208258 001	Apr 03, 2017

TAZORAC

AB	+! ALLERGAN	0.05%	N021184 001	Sep 29, 2000
AB	+!	0.1%	N021184 002	Sep 29, 2000

GEL; TOPICAL

TAZAROTENE

AB	COSETTE	0.05%	A215433 001	Sep 13, 2022
AB		0.1%	A214136 001	Sep 13, 2022
AB	PADAGIS ISRAEL	0.05%	A213079 001	Apr 25, 2023
AB		0.1%	A213079 002	Apr 25, 2023
AB	SOLARIS PHARMA CORP	0.05%	A213644 001	Mar 20, 2023
AB		0.1%	A213644 002	Mar 20, 2023

TAZORAC

AB	+! ALLERGAN	0.05%	N020600 001	Jun 13, 1997
AB	+!	0.1%	N020600 002	Jun 13, 1997

LOTION; TOPICAL

ARAZLO

+! BAUSCH 0.045%

N211882 001 Dec 18, 2019

TAZEMETOSTAT HYDROBROMIDE

TABLET; ORAL

TAZVERIK

+! EPIZYME INC EQ 200MG BASE

N211723 001 Jan 23, 2020

TECHNETIUM TC-99M BICISATE KIT

INJECTABLE; INJECTION

NEUROLITE

+! LANTHEUS MEDCL N/A

N020256 001 Nov 23, 1994

TECHNETIUM TC-99M EXAMETAZIME KIT

INJECTABLE; INJECTION

CERETEC

+! GE HEALTHCARE N/A

N019829 001 Dec 30, 1988

POWDER; INTRAVENOUS

DRAX EXAMETAZIME

JUBILANT N/A

N208870 001 Aug 17, 2017

TECHNETIUM TC-99M LABELED CARBON

AEROSOL; INHALATION

TECHNEGAS KIT

+! CYCLOMEDICA N/A

N022335 001 Sep 29, 2023

TECHNETIUM TC-99M MEBROFENIN KIT

INJECTABLE; INJECTION

CHOLETEC

AP	+! BRACCO	N/A	N018963 001	Jan 21, 1987
-----------	-----------	------------	--------------------	--------------

TECHNETIUM TC-99M MEBROFENIN

AP	SUN PHARM INDS INC	N/A	A078242 001	Jan 29, 2008
-----------	--------------------	------------	--------------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

TECHNETIUM TC-99M MEDRONATE

INJECTABLE; INJECTION

DRAXIMAGE MDP-25

+! JUBILANT N/A

N018035 002 Feb 27, 2004

TECHNETIUM TC-99M MEDRONATE KIT

INJECTABLE; INJECTION

CIS-MDP**AP** SUN PHARM INDS INC **N/A****N018124 001****TECHNETIUM TC-99M MEDRONATE KIT****AP** CARDINAL HEALTH 414 **N/A****N018107 001**TECHNETIUM TC-99M MERTIATIDE KIT

INJECTABLE; INJECTION

TECHNESCAN MAG3**AP** +! CURIUM **N/A****N019882 001** Jun 15, 1990**TECHNETIUM TC99M MERTIATIDE KIT****AP** MEDI-RADIOPHARMA **N/A****A206489 001** Feb 06, 2020**AP** SUN PHARM INDS INC **N/A****A208994 001** Jul 12, 2019

POWDER; INTRAVENOUS

TECHNETIUM TC 99M MERTIATIDE KIT

+! JUBILANT DRAXIMAGE N/A

N216820 001 Jan 30, 2023

TECHNETIUM TC-99M OXIDRONATE KIT

INJECTABLE; INJECTION

TECHNESCAN

+! CURIUM N/A

N018321 001

TECHNETIUM TC-99M PENTETATE KIT

INJECTABLE; INJECTION

DRAXIMAGE DTPA

+! JUBILANT N/A

N018511 001 Dec 29, 1989

TECHNETIUM TC-99M PYROPHOSPHATE KIT

INJECTABLE; INJECTION

CIS-PYRO**AP** SUN PHARM INDS INC **N/A****N019039 001** Jun 30, 1987**TECHNESCAN PYP KIT****AP** CURIUM **N/A****N017538 001**TECHNETIUM TC-99M RED BLOOD CELL KIT

INJECTABLE; INJECTION

ULTRATAG

+! CURIUM N/A

N019981 001 Jun 10, 1991

TECHNETIUM TC-99M SESTAMIBI KIT

INJECTABLE; INJECTION

CARDIOLITE**AP** +! LANTHEUS MEDCL **N/A****N019785 001** Dec 21, 1990**TECHNETIUM TC 99M SESTAMIBI****AP** CARDINAL HEALTH 414 **N/A****A078809 001** Apr 28, 2009**AP** CURIUM **N/A****A078098 001** Sep 22, 2008**AP** JUBILANT DRAXIMAGE **N/A****A078806 001** Apr 29, 2009**AP** SUN PHARM INDS INC **10-30mCi****A079157 001** Jul 10, 2009TECHNETIUM TC-99M SODIUM PERTECHNETATE GENERATOR

SOLUTION; INTRAVENOUS

TECHNELITE

+! LANTHEUS MEDCL 1-20 CI/GENERATOR

N017771 002 Feb 12, 2014

ULTRA-TECHNEKOW FM

+! CURIUM 1-19 CI/GENERATOR

N017243 003 Feb 18, 2014

TECHNETIUM TC-99M SUCCIMER

POWDER; INTRAVENOUS

NEPHROSCAN

+! THERAGNOSTICS N/A

N214993 001 Feb 18, 2022

TECHNETIUM TC-99M SULFUR COLLOID KIT

SOLUTION; INJECTION, ORAL

AN-SULFUR COLLOID**AP** +! SUN PHARM INDS INC **N/A****N017858 001****TECHNETIUM TC-99M SULFUR COLLOID KIT****AP** JUBILANT **N/A****A213516 001** Nov 09, 2023

PRESCRIPTION DRUG PRODUCT LISTTECHNETIUM TC-99M TETROFOSMIN KIT

INJECTABLE; INJECTION

MYOVIEW

+! MEDI-PHYSICS N/A N020372 001 Feb 09, 1996

MYOVIEW 30ML

+! MEDI-PHYSICS N/A N020372 002 Jul 07, 2005

TECHNETIUM TC-99M TILMANOCEPT

INJECTABLE; INJECTION

LYMPHOSEEK KIT

+! CARDINAL HEALTH 414 N/A N202207 001 Mar 13, 2013

TECOVIRIMAT

CAPSULE; ORAL

TPOXX

+! SIGA TECHNOLOGIES 200MG N208627 001 Jul 13, 2018

SOLUTION; INTRAVENOUS

TPOXX

+! SIGA TECHNOLOGIES 200MG/20ML (10MG/ML) N214518 001 May 18, 2022

TEDIZOLID PHOSPHATE

POWDER; INTRAVENOUS

SIVEXTRO

+! CUBIST PHARMS LLC 200MG/VIAL N205436 001 Jun 20, 2014

TABLET; ORAL

SIVEXTRO

+! CUBIST PHARMS LLC 200MG N205435 001 Jun 20, 2014

TEDUGLUTIDE

POWDER; SUBCUTANEOUS

GATTEX KIT

+! TAKEDA PHARMS USA 5MG/VIAL N203441 001 Dec 21, 2012

TELAVANCIN HYDROCHLORIDE

POWDER; INTRAVENOUS

VIBATIV

+! CUMBERLAND EQ 750MG BASE/VIAL N022110 002 Sep 11, 2009

TELMISARTAN

TABLET; ORAL

MICARDIS**AB** + BOEHRINGER 20MG **N020850 003** Apr 04, 2000
INGELHEIM**AB** + 40MG **N020850 001** Nov 10, 1998**AB** +! 80MG **N020850 002** Nov 10, 1998TELMISARTAN**AB** ALEMBIC 20MG **A202130 001** Jul 07, 2014**AB** 40MG **A202130 002** Jul 07, 2014**AB** 80MG **A202130 003** Jul 07, 2014**AB** AMNEAL PHARMS 20MG **A204415 001** Sep 08, 2015**AB** 40MG **A204415 002** Sep 08, 2015**AB** 80MG **A204415 003** Sep 08, 2015**AB** AUROBINDO PHARMA 20MG **A206511 001** Sep 03, 2015**AB** 40MG **A206511 002** Sep 03, 2015**AB** 80MG **A206511 003** Sep 03, 2015**AB** CADILA PHARMS LTD 20MG **A208605 001** Jul 25, 2017**AB** 40MG **A208605 002** Jul 25, 2017**AB** 80MG **A208605 003** Jul 25, 2017**AB** CHARTWELL RX 20MG **A078710 001** Jan 08, 2014**AB** 40MG **A078710 002** Jan 08, 2014**AB** 80MG **A078710 003** Jan 08, 2014**AB** GLENMARK PHARMS LTD 20MG **A090032 001** Jul 07, 2014**AB** 40MG **A090032 002** Jul 07, 2014**AB** 80MG **A090032 003** Jul 07, 2014**AB** HETERO LABS LTD V 20MG **A205901 001** Apr 22, 2016**AB** 40MG **A205901 002** Apr 22, 2016**AB** 80MG **A205901 003** Apr 22, 2016**AB** INVENTIA 20MG **A205150 001** Oct 30, 2015**AB** 40MG **A205150 002** Oct 30, 2015**AB** 80MG **A205150 003** Oct 30, 2015**AB** MANKIND PHARMA 20MG **A218157 001** Oct 15, 2024**AB** 40MG **A218157 002** Oct 15, 2024**AB** 80MG **A218157 003** Oct 15, 2024**AB** MICRO LABS 20MG **A207016 001** Oct 03, 2017**AB** 40MG **A207016 002** Oct 03, 2017**AB** 80MG **A207016 003** Oct 03, 2017

PRESCRIPTION DRUG PRODUCT LIST

TELMISARTAN

TABLET; ORAL

TELMISARTAN

<u>AB</u>	MYLAN	<u>20MG</u>	<u>A202397 001</u>	Jul 07, 2014
<u>AB</u>		<u>40MG</u>	<u>A202397 002</u>	Jul 07, 2014
<u>AB</u>		<u>80MG</u>	<u>A202397 003</u>	Jul 07, 2014
<u>AB</u>	PRINSTON INC	<u>20MG</u>	<u>A207882 001</u>	May 03, 2017
<u>AB</u>		<u>40MG</u>	<u>A207882 002</u>	May 03, 2017
<u>AB</u>		<u>80MG</u>	<u>A207882 003</u>	May 03, 2017
<u>AB</u>	SANDOZ	<u>20MG</u>	<u>A203867 001</u>	Nov 03, 2014
<u>AB</u>		<u>40MG</u>	<u>A203867 002</u>	Nov 03, 2014
<u>AB</u>		<u>80MG</u>	<u>A203867 003</u>	Nov 03, 2014
<u>AB</u>	ZYDUS PHARMS	<u>20MG</u>	<u>A203325 001</u>	Aug 26, 2014
<u>AB</u>		<u>40MG</u>	<u>A203325 002</u>	Aug 26, 2014
<u>AB</u>		<u>80MG</u>	<u>A203325 003</u>	Aug 26, 2014

TELOTRISTAT ETIPRATE

TABLET; ORAL

XERMELO

+! TERSERA

EQ 250MG BASE

N208794 001 Feb 28, 2017

TEMAZEPAM

CAPSULE; ORAL

RESTORIL

<u>AB</u>	+ SPECGX LLC	<u>7.5MG</u>	<u>N018163 003</u>	Oct 25, 1991
<u>AB</u>	+	<u>15MG</u>	<u>N018163 001</u>	
<u>AB</u>	+	<u>22.5MG</u>	<u>N018163 004</u>	Nov 02, 2004
<u>AB</u>	+!	<u>30MG</u>	<u>N018163 002</u>	

TEMAZEPAM

<u>AB</u>	ACTAVIS ELIZABETH	<u>15MG</u>	<u>A071620 002</u>	Aug 07, 1987
<u>AB</u>		<u>30MG</u>	<u>A071620 001</u>	Aug 07, 1987
<u>AB</u>	ALEMBIC	<u>7.5MG</u>	<u>A211542 001</u>	Nov 23, 2018
<u>AB</u>		<u>15MG</u>	<u>A211542 002</u>	Nov 23, 2018
<u>AB</u>		<u>22.5MG</u>	<u>A211542 003</u>	Nov 23, 2018
<u>AB</u>		<u>30MG</u>	<u>A211542 004</u>	Nov 23, 2018
<u>AB</u>	ALKEM LABS LTD	<u>7.5MG</u>	<u>A217875 001</u>	Aug 24, 2023
<u>AB</u>		<u>15MG</u>	<u>A217875 002</u>	Aug 24, 2023
<u>AB</u>		<u>22.5MG</u>	<u>A217875 003</u>	Aug 24, 2023
<u>AB</u>		<u>30MG</u>	<u>A217875 004</u>	Aug 24, 2023
<u>AB</u>	CHARTWELL RX	<u>15MG</u>	<u>A071427 001</u>	Jan 12, 1988
<u>AB</u>		<u>30MG</u>	<u>A071428 001</u>	Jan 12, 1988
<u>AB</u>	NOVEL LABS INC	<u>7.5MG</u>	<u>A071457 002</u>	Jun 22, 2012
<u>AB</u>		<u>15MG</u>	<u>A071456 001</u>	Apr 21, 1987
<u>AB</u>		<u>22.5MG</u>	<u>A071457 003</u>	Jun 22, 2012
<u>AB</u>		<u>30MG</u>	<u>A071457 001</u>	Apr 21, 1987
<u>AB</u>	PRINSTON INC	<u>7.5MG</u>	<u>A201781 001</u>	Jun 04, 2015
<u>AB</u>		<u>15MG</u>	<u>A201781 002</u>	Jun 04, 2015
<u>AB</u>		<u>22.5MG</u>	<u>A201781 003</u>	Jun 04, 2015
<u>AB</u>		<u>30MG</u>	<u>A201781 004</u>	Jun 04, 2015
<u>AB</u>	SUN PHARM INDUSTRIES	<u>7.5MG</u>	<u>A078581 001</u>	Sep 08, 2009

TEMOZOLOMIDE

CAPSULE; ORAL

TEMOZOLOMIDE

<u>AB</u>	ACCORD HLTHCARE	<u>5MG</u>	<u>A201528 001</u>	Feb 27, 2017
<u>AB</u>		<u>20MG</u>	<u>A201528 002</u>	Feb 27, 2017
<u>AB</u>		<u>100MG</u>	<u>A201528 003</u>	Feb 27, 2017
<u>AB</u>		<u>140MG</u>	<u>A201528 004</u>	Feb 27, 2017
<u>AB</u>		<u>180MG</u>	<u>A201528 005</u>	Feb 27, 2017
<u>AB</u>		<u>250MG</u>	<u>A201528 006</u>	Feb 27, 2017
<u>AB</u>	AMNEAL PHARMS	<u>5MG</u>	<u>A203691 001</u>	May 08, 2015
<u>AB</u>		<u>20MG</u>	<u>A203691 002</u>	May 08, 2015
<u>AB</u>		<u>100MG</u>	<u>A203691 003</u>	May 08, 2015
<u>AB</u>		<u>140MG</u>	<u>A203691 004</u>	May 08, 2015
<u>AB</u>		<u>180MG</u>	<u>A203691 005</u>	May 08, 2015
<u>AB</u>	!	<u>250MG</u>	<u>A203691 006</u>	May 08, 2015
<u>AB</u>	CHARTWELL	<u>5MG</u>	<u>A206413 001</u>	Apr 12, 2016
<u>AB</u>		<u>20MG</u>	<u>A206413 002</u>	Apr 12, 2016
<u>AB</u>		<u>100MG</u>	<u>A206413 003</u>	Apr 12, 2016
<u>AB</u>		<u>140MG</u>	<u>A206413 004</u>	Apr 12, 2016
<u>AB</u>		<u>180MG</u>	<u>A206413 005</u>	Apr 12, 2016
<u>AB</u>		<u>250MG</u>	<u>A206413 006</u>	Apr 12, 2016
<u>AB</u>	CHEMI SPA	<u>5MG</u>	<u>A204639 001</u>	Nov 23, 2016

PRESCRIPTION DRUG PRODUCT LIST

TEMOZOLOMIDE

CAPSULE; ORAL

TEMOZOLOMIDE

<u>AB</u>		<u>20MG</u>	<u>A204639 002</u>	Nov 23, 2016
<u>AB</u>		<u>100MG</u>	<u>A204639 003</u>	Nov 23, 2016
<u>AB</u>		<u>140MG</u>	<u>A204639 004</u>	Nov 23, 2016
<u>AB</u>		<u>180MG</u>	<u>A204639 005</u>	Nov 23, 2016
<u>AB</u>		<u>250MG</u>	<u>A204639 006</u>	Nov 23, 2016
<u>AB</u>	DEVA HOLDING AS	<u>5MG</u>	<u>A207658 001</u>	Apr 26, 2017
<u>AB</u>		<u>20MG</u>	<u>A207658 002</u>	Apr 26, 2017
<u>AB</u>		<u>100MG</u>	<u>A207658 003</u>	Apr 26, 2017
<u>AB</u>		<u>140MG</u>	<u>A207658 004</u>	Apr 26, 2017
<u>AB</u>		<u>180MG</u>	<u>A207658 005</u>	Apr 26, 2017
<u>AB</u>		<u>250MG</u>	<u>A207658 006</u>	Apr 26, 2017
<u>AB</u>	HETERO LABS LTD V	<u>5MG</u>	<u>A210030 001</u>	Aug 23, 2024
<u>AB</u>		<u>20MG</u>	<u>A210030 002</u>	Aug 23, 2024
<u>AB</u>		<u>100MG</u>	<u>A210030 003</u>	Aug 23, 2024
<u>AB</u>		<u>140MG</u>	<u>A210030 004</u>	Aug 23, 2024
<u>AB</u>		<u>180MG</u>	<u>A210030 005</u>	Aug 23, 2024
<u>AB</u>		<u>250MG</u>	<u>A210030 006</u>	Aug 23, 2024
<u>AB</u>	RISING	<u>5MG</u>	<u>A206309 001</u>	Apr 27, 2016
<u>AB</u>		<u>20MG</u>	<u>A206309 002</u>	Apr 27, 2016
<u>AB</u>		<u>100MG</u>	<u>A206309 003</u>	Apr 27, 2016
<u>AB</u>		<u>140MG</u>	<u>A206309 004</u>	Apr 27, 2016
<u>AB</u>		<u>180MG</u>	<u>A206309 005</u>	Apr 27, 2016
<u>AB</u>		<u>250MG</u>	<u>A206309 006</u>	Apr 27, 2016
<u>AB</u>	SUN PHARM	<u>5MG</u>	<u>A201742 001</u>	Feb 12, 2014
<u>AB</u>		<u>20MG</u>	<u>A201742 002</u>	Feb 12, 2014
<u>AB</u>		<u>100MG</u>	<u>A201742 003</u>	Feb 12, 2014
<u>AB</u>		<u>140MG</u>	<u>A201742 004</u>	Feb 12, 2014
<u>AB</u>		<u>180MG</u>	<u>A201742 005</u>	Feb 12, 2014
<u>AB</u>		<u>250MG</u>	<u>A201742 006</u>	Feb 12, 2014
<u>AB</u>	ZYDUS PHARMS	<u>5MG</u>	<u>A206750 001</u>	Jul 31, 2017
<u>AB</u>		<u>20MG</u>	<u>A206750 002</u>	Jul 31, 2017
<u>AB</u>		<u>100MG</u>	<u>A206750 003</u>	Jul 31, 2017
<u>AB</u>		<u>140MG</u>	<u>A206750 004</u>	Jul 31, 2017
<u>AB</u>		<u>180MG</u>	<u>A206750 005</u>	Jul 31, 2017
<u>AB</u>		<u>250MG</u>	<u>A206750 006</u>	Jul 31, 2017

POWDER; INTRAVENOUS

TEMODAR

+! MERCK SHARP DOHME 100MG/VIAL N022277 001 Feb 27, 2009

TEMSIROLIMUS

SOLUTION; INTRAVENOUS

TEMSIROLIMUS

<u>AP</u>	ACCORD HLTHCARE	<u>25MG/ML (25MG/ML)</u>	<u>A203153 001</u>	Jul 30, 2018
<u>AP</u>	GLAND PHARMA LTD	<u>25MG/ML (25MG/ML)</u>	<u>A207383 001</u>	Aug 16, 2019
<u>AP</u>	+! PF PRISM CV	<u>25MG/ML (25MG/ML)</u>	<u>N022088 001</u>	May 30, 2007

TENAPANOR HYDROCHLORIDE

TABLET; ORAL

IBSRELA

+! ARDELYX INC EQ 50MG BASE N211801 001 Sep 12, 2019

XPHOZAH

+ ARDELYX INC EQ 20MG BASE N213931 002 Oct 17, 2023

+! EQ 30MG BASE N213931 003 Oct 17, 2023

TENOFOVIR ALAFENAMIDE FUMARATE

TABLET; ORAL

TENOFOVIR ALAFENAMIDEAB LUPIN LTD EQ 25MG BASE A214226 001 Mar 30, 2023VEMLIDYAB +! GILEAD SCIENCES INC EQ 25MG BASE N208464 001 Nov 10, 2016TENOFOVIR DISOPROXIL FUMARATE

POWDER; ORAL

VIREAD

+! GILEAD SCIENCES INC 40MG/SCOOPFUL N022577 001 Jan 18, 2012

TABLET; ORAL

TENOFOVIR DISOPROXIL FUMARATE

<u>AB</u>	AUROBINDO PHARMA	<u>150MG</u>	<u>A090647 001</u>	Jan 26, 2018
<u>AB</u>		<u>200MG</u>	<u>A090647 002</u>	Jan 26, 2018
<u>AB</u>		<u>250MG</u>	<u>A090647 003</u>	Jan 26, 2018

PRESCRIPTION DRUG PRODUCT LIST

TENOFOVIR DISOPROXIL FUMARATE

TABLET; ORAL

TENOFOVIR DISOPROXIL FUMARATE

<u>AB</u>		<u>300MG</u>	<u>A090647</u>	<u>004</u>	Jan 26, 2018
<u>AB</u>	CHARTWELL	<u>300MG</u>	<u>A206481</u>	<u>001</u>	Jul 26, 2018
<u>AB</u>	CIPLA	<u>300MG</u>	<u>A078800</u>	<u>001</u>	Jan 26, 2018
<u>AB</u>	HETERO LABS LTD III	<u>300MG</u>	<u>A090636</u>	<u>001</u>	Jan 26, 2018
<u>AB</u>	MACLEODS PHARMS LTD	<u>300MG</u>	<u>A203232</u>	<u>001</u>	Jan 26, 2018
<u>AB</u>	NORVIUM BIOSCIENCE	<u>300MG</u>	<u>A079071</u>	<u>001</u>	Feb 29, 2024
<u>AB</u>	QILU	<u>200MG</u>	<u>A209498</u>	<u>001</u>	Mar 02, 2018
<u>AB</u>		<u>250MG</u>	<u>A209498</u>	<u>002</u>	Mar 02, 2018
<u>AB</u>		<u>300MG</u>	<u>A209498</u>	<u>003</u>	Mar 02, 2018
<u>AB</u>	REYOUNG	<u>300MG</u>	<u>A211337</u>	<u>001</u>	Mar 02, 2023
<u>AB</u>	STRIDES PHARMA	<u>300MG</u>	<u>A090742</u>	<u>001</u>	Jan 26, 2018
<u>AB</u>	TEVA PHARMS USA	<u>150MG</u>	<u>A091612</u>	<u>002</u>	Jan 26, 2018
<u>AB</u>		<u>200MG</u>	<u>A091612</u>	<u>003</u>	Jan 26, 2018
<u>AB</u>		<u>250MG</u>	<u>A091612</u>	<u>004</u>	Jan 26, 2018
<u>AB</u>		<u>300MG</u>	<u>A091612</u>	<u>001</u>	Mar 18, 2015
<u>VIREAD</u>					
<u>AB</u>	+ GILEAD SCIENCES INC	<u>150MG</u>	<u>N021356</u>	<u>002</u>	Jan 18, 2012
<u>AB</u>	+	<u>200MG</u>	<u>N021356</u>	<u>003</u>	Jan 18, 2012
<u>AB</u>	+	<u>250MG</u>	<u>N021356</u>	<u>004</u>	Jan 18, 2012
<u>AB</u>	+	<u>300MG</u>	<u>N021356</u>	<u>001</u>	Oct 26, 2001

TEPOTINIB HYDROCHLORIDE

TABLET; ORAL

TEPMETKO

+! EMD SERONO INC EQ 225MG BASE N214096 001 Feb 03, 2021

TERAZOSIN HYDROCHLORIDE

CAPSULE; ORAL

TERAZOSIN HYDROCHLORIDE

<u>AB</u>	APNAR PHARMA LP	<u>EQ 1MG BASE</u>	<u>A074823</u>	<u>001</u>	Mar 30, 1998
<u>AB</u>		<u>EQ 2MG BASE</u>	<u>A074823</u>	<u>002</u>	Mar 30, 1998
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A074823</u>	<u>003</u>	Mar 30, 1998
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A074823</u>	<u>004</u>	Mar 30, 1998
<u>AB</u>	BIONPHARMA	<u>EQ 1MG BASE</u>	<u>A075667</u>	<u>001</u>	Jul 28, 2000
<u>AB</u>		<u>EQ 2MG BASE</u>	<u>A075667</u>	<u>002</u>	Jul 28, 2000
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A075667</u>	<u>003</u>	Jul 28, 2000
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A075667</u>	<u>004</u>	Jul 28, 2000
<u>AB</u>	HERITAGE PHARMA	<u>EQ 1MG BASE</u>	<u>A075614</u>	<u>002</u>	Jan 30, 2001
<u>AB</u>	AVET	<u>EQ 2MG BASE</u>	<u>A075614</u>	<u>001</u>	Jan 30, 2001
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A075614</u>	<u>003</u>	Jan 30, 2001
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A075614</u>	<u>004</u>	Jan 30, 2001
<u>AB</u>	! JUBILANT CADISTA	<u>EQ 1MG BASE</u>	<u>A075317</u>	<u>001</u>	Dec 20, 2004
<u>AB</u>		<u>EQ 2MG BASE</u>	<u>A075317</u>	<u>002</u>	Dec 20, 2004
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A075317</u>	<u>003</u>	Dec 20, 2004
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A075317</u>	<u>004</u>	Dec 20, 2004

SOLUTION; ORAL

TEZRULY

+! NOVITIUM PHARMA EQ 1MG BASE/ML N218139 001 Jul 29, 2024

TERBINAFINE HYDROCHLORIDE

TABLET; ORAL

TERBINAFINE HYDROCHLORIDE

<u>AB</u>	! AUROBINDO PHARMA	<u>EQ 250MG BASE</u>	<u>A078297</u>	<u>001</u>	Jul 02, 2007
<u>AB</u>	BRECKENRIDGE PHARM	<u>EQ 250MG BASE</u>	<u>A077714</u>	<u>001</u>	Jun 04, 2010
<u>AB</u>	CHARTWELL	<u>EQ 250MG BASE</u>	<u>A078199</u>	<u>001</u>	Jul 02, 2007
<u>AB</u>	DR REDDYS LABS INC	<u>EQ 250MG BASE</u>	<u>A076390</u>	<u>001</u>	Jul 02, 2007
<u>AB</u>	EMED MEDCL	<u>EQ 250MG BASE</u>	<u>A077919</u>	<u>001</u>	Jul 02, 2007
<u>AB</u>	GLENMARK PHARMS LTD	<u>EQ 250MG BASE</u>	<u>A078157</u>	<u>001</u>	Jul 02, 2007
<u>AB</u>	INVAGEN PHARMS	<u>EQ 250MG BASE</u>	<u>A077533</u>	<u>001</u>	Jul 02, 2007
<u>AB</u>	ORBION PHARMS	<u>EQ 250MG BASE</u>	<u>A078163</u>	<u>001</u>	Jul 02, 2007

TERBUTALINE SULFATE

INJECTABLE; INJECTION

TERBUTALINE SULFATE

<u>AP</u>	AREVA PHARMS	<u>1MG/ML</u>	<u>A200122</u>	<u>001</u>	Nov 08, 2013
<u>AP</u>	FRESENIUS KABI USA	<u>1MG/ML</u>	<u>A076887</u>	<u>001</u>	May 26, 2004
<u>AP</u>	! HIKMA FARMACEUTICA	<u>1MG/ML</u>	<u>A078630</u>	<u>001</u>	May 20, 2009

TABLET; ORAL

BRETHINE

<u>AB</u>	+ ANI PHARMS	<u>2.5MG</u>	<u>N017849</u>	<u>001</u>	
<u>AB</u>	+	<u>5MG</u>	<u>N017849</u>	<u>002</u>	

PRESCRIPTION DRUG PRODUCT LIST

TERBUTALINE SULFATE

TABLET; ORAL

TERBUTALINE SULFATE

<u>AB</u>	IMPAX LABS	<u>2.5MG</u>	<u>A075877 001</u>	Jun 26, 2001
<u>AB</u>		<u>5MG</u>	<u>A075877 002</u>	Jun 26, 2001
<u>AB</u>	LANNETT CO INC	<u>2.5MG</u>	<u>A077152 001</u>	Mar 25, 2005
<u>AB</u>	!	<u>5MG</u>	<u>A077152 002</u>	Mar 25, 2005
<u>AB</u>	TWI PHARMS	<u>2.5MG</u>	<u>A211832 001</u>	Jun 19, 2020
<u>AB</u>		<u>5MG</u>	<u>A211832 002</u>	Jun 19, 2020

TERCONAZOLE

CREAM; VAGINAL

TERCONAZOLE

<u>AB</u>	FOUGERA PHARMS	<u>0.4%</u>	<u>A076712 001</u>	Feb 18, 2005
<u>AB</u>	!	<u>0.4%</u>	<u>A076043 001</u>	Jan 19, 2005
BX	+!	FOUGERA PHARMS INC	N021735 001	Oct 01, 2004
BX	!	TARO	A075953 001	Apr 06, 2004

SUPPOSITORY; VAGINAL

TERCONAZOLE

<u>AB</u>	!	PADAGIS ISRAEL	<u>80MG</u>	<u>A077149 001</u>	Mar 17, 2006
<u>AB</u>		TARO	<u>80MG</u>	<u>A077553 001</u>	Mar 09, 2007

TERIFLUNOMIDE

TABLET; ORAL

AUBAGIO

<u>AB</u>	+	SANOFI AVENTIS US	<u>7MG</u>	<u>N202992 001</u>	Sep 12, 2012
<u>AB</u>	+!		<u>14MG</u>	<u>N202992 002</u>	Sep 12, 2012

TERIFLUNOMIDE

<u>AB</u>		ACCORD HLTHCARE	<u>7MG</u>	<u>A209690 001</u>	Jan 07, 2019
<u>AB</u>			<u>14MG</u>	<u>A209690 002</u>	Jan 07, 2019
<u>AB</u>		ALEMBIC	<u>7MG</u>	<u>A209572 001</u>	Apr 19, 2019
<u>AB</u>			<u>14MG</u>	<u>A209572 002</u>	Apr 19, 2019
<u>AB</u>		APOTEX	<u>7MG</u>	<u>A209601 001</u>	Nov 02, 2018
<u>AB</u>			<u>14MG</u>	<u>A209601 002</u>	Nov 02, 2018
<u>AB</u>		AUROBINDO PHARMA	<u>7MG</u>	<u>A209638 001</u>	Oct 26, 2018
<u>AB</u>			<u>14MG</u>	<u>A209638 002</u>	Oct 26, 2018
<u>AB</u>		BIOCON PHARMA	<u>7MG</u>	<u>A209639 001</u>	Mar 13, 2023
<u>AB</u>			<u>14MG</u>	<u>A209639 002</u>	Mar 13, 2023
<u>AB</u>		GLENMARK SPECLT	<u>7MG</u>	<u>A209663 001</u>	Nov 15, 2018
<u>AB</u>			<u>14MG</u>	<u>A209663 002</u>	Nov 15, 2018
<u>AB</u>		HETERO LABS LTD V	<u>7MG</u>	<u>A209598 001</u>	Mar 13, 2023
<u>AB</u>			<u>14MG</u>	<u>A209598 002</u>	Mar 13, 2023
<u>AB</u>		MSN	<u>7MG</u>	<u>A209623 001</u>	Apr 24, 2019
<u>AB</u>			<u>14MG</u>	<u>A209623 002</u>	Apr 24, 2019
<u>AB</u>		NATCO	<u>7MG</u>	<u>A209555 001</u>	May 15, 2023
<u>AB</u>			<u>14MG</u>	<u>A209555 002</u>	May 15, 2023
<u>AB</u>		SANDOZ	<u>7MG</u>	<u>A209710 001</u>	Jan 03, 2019
<u>AB</u>			<u>14MG</u>	<u>A209710 002</u>	Jan 03, 2019
<u>AB</u>		SOLA PHARMS	<u>7MG</u>	<u>A209677 002</u>	Sep 28, 2023
<u>AB</u>			<u>14MG</u>	<u>A209677 001</u>	Jun 17, 2020
<u>AB</u>		TEVA PHARMS USA	<u>7MG</u>	<u>A209700 001</u>	Sep 04, 2018
<u>AB</u>			<u>14MG</u>	<u>A209700 002</u>	Sep 04, 2018
<u>AB</u>		ZYDUS PHARMS	<u>7MG</u>	<u>A209668 001</u>	Nov 30, 2018
<u>AB</u>			<u>14MG</u>	<u>A209668 002</u>	Nov 30, 2018

TERIPARATIDE

SOLUTION; SUBCUTANEOUS

FORTEO

<u>AP</u>	+!	LILLY	<u>0.6MG/2.4ML (0.25MG/ML)</u>	<u>N021318 002</u>	Jun 25, 2008
-----------	----	-------	--------------------------------	--------------------	--------------

TERIPARATIDE

<u>AP</u>		APOTEX	<u>0.6MG/2.4ML (0.25MG/ML)</u>	<u>A211097 001</u>	Nov 16, 2023	
<u>AP</u>		TEVA PHARMS USA	<u>0.6MG/2.4ML (0.25MG/ML)</u>	<u>A208569 001</u>	Nov 16, 2023	
		BONSITY				
		ALVOGEN	0.62MG/2.48ML (0.25MG/ML)	N211939 001	Oct 04, 2019	
		TERIPARATIDE				
		+!	ALMAJECT	0.6MG/2.4ML (0.25MG/ML)	N218771 001	Jun 04, 2024

TERLIPRESSIN ACETATE

POWDER; INTRAVENOUS

TERLIVAZ

		+!	MALLINCKRODT IRELAND	EQ 0.85MG BASE/VIAL	N022231 001	Sep 14, 2022
--	--	----	----------------------	---------------------	-------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

TESTOSTERONE

GEL; TRANSDERMAL

ANDROGEL

<u>AB1</u>	+	BESINS HLTHCARE	<u>25MG/2.5GM PACKET</u>	<u>N021015</u>	<u>001</u>	Feb 28, 2000
<u>AB1</u>	+		<u>50MG/5GM PACKET</u>	<u>N021015</u>	<u>002</u>	Feb 28, 2000

TESTOSTERONE

<u>AB1</u>		ACTAVIS LABS UT INC	<u>25MG/2.5GM PACKET</u>	<u>A076737</u>	<u>001</u>	Jan 27, 2006
<u>AB1</u>	!		<u>50MG/5GM PACKET</u>	<u>A076737</u>	<u>002</u>	Jan 27, 2006
<u>AB1</u>		ENCUBE	<u>25MG/2.5GM PACKET</u>	<u>A212984</u>	<u>003</u>	Jul 11, 2024
<u>AB1</u>			<u>50MG/5GM PACKET</u>	<u>A212984</u>	<u>001</u>	Nov 09, 2021
<u>AB1</u>		STRIDES PHARMA	<u>25MG/2.5GM PACKET</u>	<u>A076744</u>	<u>001</u>	May 23, 2007
<u>AB1</u>			<u>50MG/5GM PACKET</u>	<u>A076744</u>	<u>002</u>	May 23, 2007

ANDROGEL

<u>AB2</u>	+	BESINS HLTHCARE	<u>1.62% (20.25MG/1.25GM PACKET)</u>	<u>N022309</u>	<u>002</u>	Sep 07, 2012
<u>AB2</u>	+		<u>1.62% (40.5MG/2.5GM PACKET)</u>	<u>N022309</u>	<u>003</u>	Sep 07, 2012

TESTIM

<u>AB2</u>	+	ENDO OPERATIONS	<u>50MG/5GM PACKET</u>	<u>N021454</u>	<u>001</u>	Oct 31, 2002
------------	---	-----------------	------------------------	----------------	------------	--------------

TESTOSTERONE

<u>AB2</u>		ACTAVIS LABS UT INC	<u>1.62% (20.25MG/1.25GM PACKET)</u>	<u>A204570</u>	<u>002</u>	Jul 17, 2020
<u>AB2</u>			<u>1.62% (40.5MG/2.5GM PACKET)</u>	<u>A204570</u>	<u>003</u>	Jul 17, 2020
<u>AB2</u>			<u>50MG/5GM PACKET</u>	<u>A091073</u>	<u>001</u>	Sep 18, 2017
<u>AB2</u>		PADAGIS ISRAEL	<u>1.62% (20.25MG/1.25GM PACKET)</u>	<u>A205781</u>	<u>001</u>	Jul 12, 2017
<u>AB2</u>	!		<u>1.62% (40.5MG/2.5GM PACKET)</u>	<u>A205781</u>	<u>002</u>	Jul 12, 2017

VOGELXO

<u>AB2</u>		UPSHER SMITH LABS	<u>50MG/5GM PACKET</u>	<u>N204399</u>	<u>002</u>	Jun 04, 2014
------------	--	-------------------	------------------------	----------------	------------	--------------

GEL, METERED; NASAL

NATESTO

+!

ACERUS

5.5MG/0.122GM ACTUATION

N205488 001 May 28, 2014

GEL, METERED; TRANSDERMAL

ANDROGEL

<u>AB</u>	+	BESINS HLTHCARE	<u>1.62% (20.25MG/1.25GM ACTUATION)</u>	<u>N022309</u>	<u>001</u>	Apr 29, 2011
-----------	---	-----------------	---	----------------	------------	--------------

TESTOSTERONE

<u>AB</u>		ACTAVIS LABS UT INC	<u>1.62% (20.25MG/1.25GM ACTUATION)</u>	<u>A204570</u>	<u>001</u>	Apr 10, 2019
<u>AB</u>		AMNEAL	<u>1.62% (20.25MG/1.25GM ACTUATION)</u>	<u>A207373</u>	<u>001</u>	Apr 10, 2019
<u>AB</u>		ENCUBE	<u>1.62% (20.25MG/1.25GM ACTUATION)</u>	<u>A208620</u>	<u>001</u>	Apr 10, 2019
<u>AB</u>		LUPIN	<u>1.62% (20.25MG/1.25GM ACTUATION)</u>	<u>A208560</u>	<u>001</u>	Apr 10, 2019
<u>AB</u>		PADAGIS ISRAEL	<u>1.62% (20.25MG/1.25GM ACTUATION)</u>	<u>A204268</u>	<u>001</u>	Aug 04, 2015
<u>AB</u>		TWI PHARMS	<u>1.62% (20.25MG/1.25GM ACTUATION)</u>	<u>A209390</u>	<u>001</u>	Sep 23, 2019
<u>AB</u>		XIROMED	<u>1.62% (20.25MG/1.25GM ACTUATION)</u>	<u>A210835</u>	<u>001</u>	Apr 16, 2020

ANDROGEL

<u>AB1</u>	+	BESINS HLTHCARE	<u>12.5MG/1.25GM ACTUATION</u>	<u>N021015</u>	<u>003</u>	Sep 26, 2003
------------	---	-----------------	--------------------------------	----------------	------------	--------------

TESTOSTERONE

<u>AB1</u>	!	ACTAVIS LABS UT INC	<u>12.5MG/1.25GM ACTUATION</u>	<u>A076737</u>	<u>003</u>	Mar 09, 2015
<u>AB1</u>		ENCUBE	<u>12.5MG/1.25GM ACTUATION</u>	<u>A212984</u>	<u>002</u>	Feb 09, 2024

VOGELXO

BX		UPSHER SMITH LABS	12.5MG/1.25GM ACTUATION	N204399	003	Jun 04, 2014
----	--	-------------------	-------------------------	---------	-----	--------------

TESTOSTERONE

!

ACTAVIS LABS UT INC

10MG/0.5GM ACTUATION

A204571 001 Aug 05, 2015

PELLET; IMPLANTATION

TESTOPEL

!

ENDO OPERATIONS

75MG

A080911 001

SOLUTION, METERED; TRANSDERMAL

TESTOSTERONE

<u>AT</u>		ACTAVIS LABS UT INC	<u>30MG/1.5ML ACTUATION</u>	<u>A205328</u>	<u>001</u>	Aug 07, 2017
<u>AT</u>		CIPLA	<u>30MG/1.5ML ACTUATION</u>	<u>A209533</u>	<u>001</u>	Jan 29, 2018
<u>AT</u>		ENCUBE	<u>30MG/1.5ML ACTUATION</u>	<u>A212301</u>	<u>001</u>	Jan 11, 2021
<u>AT</u>		LUPIN LTD	<u>30MG/1.5ML ACTUATION</u>	<u>A208061</u>	<u>001</u>	Oct 23, 2017
<u>AT</u>	!	PADAGIS ISRAEL	<u>30MG/1.5ML ACTUATION</u>	<u>A204255</u>	<u>001</u>	Feb 28, 2017
<u>AT</u>		TWI PHARMS	<u>30MG/1.5ML ACTUATION</u>	<u>A209836</u>	<u>001</u>	Sep 03, 2021

TESTOSTERONE CYPIONATE

INJECTABLE; INJECTION

DEPO-TESTOSTERONE

<u>AO</u>	+	PFIZER	<u>100MG/ML</u>	<u>A085635</u>	<u>002</u>	
<u>AO</u>	+		<u>200MG/ML</u>	<u>A085635</u>	<u>003</u>	

TESTOSTERONE CYPIONATE

<u>AO</u>		AM REGENT	<u>200MG/ML</u>	<u>A207742</u>	<u>001</u>	Jun 16, 2017
<u>AO</u>		CIPLA	<u>100MG/ML</u>	<u>A210362</u>	<u>001</u>	Jun 19, 2018
<u>AO</u>			<u>200MG/ML</u>	<u>A210362</u>	<u>002</u>	Jun 19, 2018
<u>AO</u>		EUGIA PHARMA	<u>100MG/ML</u>	<u>A211817</u>	<u>001</u>	Oct 20, 2023
<u>AO</u>			<u>200MG/ML</u>	<u>A211817</u>	<u>002</u>	Oct 20, 2023
<u>AO</u>		HIKMA	<u>100MG/ML</u>	<u>A090387</u>	<u>001</u>	Jul 15, 2010
<u>AO</u>			<u>200MG/ML</u>	<u>A090387</u>	<u>002</u>	Jul 15, 2010

PRESCRIPTION DRUG PRODUCT LIST

TESTOSTERONE CYPIONATE

INJECTABLE; INJECTION

TESTOSTERONE CYPIONATE

<u>AO</u>	HIKMA FARMACEUTICA	<u>200MG/ML</u>	<u>A091244</u>	<u>001</u>	May 01, 2012
<u>AO</u>	PADAGIS US	<u>200MG/ML</u>	<u>A040530</u>	<u>001</u>	Jan 31, 2005
<u>AO</u>	SANDOZ	<u>100MG/ML</u>	<u>A040615</u>	<u>001</u>	Aug 10, 2006
<u>AO</u>		<u>200MG/ML</u>	<u>A040615</u>	<u>002</u>	Aug 10, 2006
<u>AO</u>	WILSHIRE PHARMS INC	<u>200MG/ML</u>	<u>A206368</u>	<u>001</u>	Apr 24, 2019
<u>AO</u>	XIROMED	<u>100MG/ML</u>	<u>A215351</u>	<u>001</u>	Apr 13, 2023
<u>AO</u>		<u>200MG/ML</u>	<u>A215351</u>	<u>002</u>	Apr 13, 2023

SOLUTION; INTRAMUSCULAR

AZMIRO

+!	AZURITY	200MG/ML (200MG/ML)	N216318	001	Jun 02, 2022
----	---------	---------------------	---------	-----	--------------

TESTOSTERONE ENANTHATE

INJECTABLE; INJECTION

TESTOSTERONE ENANTHATE

<u>AO</u>	EUGIA PHARMA	<u>200MG/ML</u>	<u>A212659</u>	<u>001</u>	Feb 28, 2024	
<u>AO</u>	!	HIKMA FARMACEUTICA	<u>200MG/ML</u>	<u>A091120</u>	<u>001</u>	Sep 18, 2012
<u>AO</u>	NEXUS	<u>200MG/ML</u>	<u>A040575</u>	<u>001</u>	Jun 14, 2006	

SOLUTION; SUBCUTANEOUS

XYOSTED (AUTOINJECTOR)

+!	ANTARES PHARMA INC	50MG/0.5ML (50MG/0.5ML)	N209863	001	Sep 28, 2018
+!		75MG/0.5ML (75MG/0.5ML)	N209863	002	Sep 28, 2018
+!		100MG/0.5ML (100MG/0.5ML)	N209863	003	Sep 28, 2018

TESTOSTERONE UNDECANOATE

CAPSULE; ORAL

JATENZO

+	TOLMAR	158MG	N206089	001	Mar 27, 2019
+		198MG	N206089	002	Mar 27, 2019
+!		237MG	N206089	003	Mar 27, 2019

KYZATREX

+	MARIUS PHARMS LLC	100MG	N213953	001	Jul 27, 2022
+		150MG	N213953	002	Jul 27, 2022
+!		200MG	N213953	003	Jul 27, 2022

TLANDO

+!	VERITY	112.5MG	N208088	001	Mar 28, 2022
----	--------	---------	---------	-----	--------------

INJECTABLE; INTRAMUSCULAR

AVEED

+!	ENDO OPERATIONS	750MG/3ML (250MG/ML)	N022219	001	Mar 05, 2014
----	-----------------	----------------------	---------	-----	--------------

TETRABENAZINE

TABLET; ORAL

TETRABENAZINE

<u>AB</u>	ACTAVIS LABS FL INC	<u>25MG</u>	<u>A206686</u>	<u>001</u>	Jul 07, 2017
<u>AB</u>	ADAPTIS	<u>12.5MG</u>	<u>A213316</u>	<u>001</u>	Jan 22, 2020
<u>AB</u>		<u>25MG</u>	<u>A213316</u>	<u>002</u>	Jan 22, 2020
<u>AB</u>	APOTEX	<u>12.5MG</u>	<u>A206093</u>	<u>001</u>	Mar 17, 2020
<u>AB</u>		<u>25MG</u>	<u>A206093</u>	<u>002</u>	Mar 17, 2020
<u>AB</u>	BIONPHARMA	<u>12.5MG</u>	<u>A208826</u>	<u>001</u>	Dec 18, 2017
<u>AB</u>		<u>25MG</u>	<u>A208826</u>	<u>002</u>	Dec 18, 2017
<u>AB</u>	CHARTWELL RX	<u>12.5MG</u>	<u>A210544</u>	<u>001</u>	Apr 20, 2018
<u>AB</u>		<u>25MG</u>	<u>A210544</u>	<u>002</u>	Apr 20, 2018
<u>AB</u>	DR REDDYS	<u>12.5MG</u>	<u>A209284</u>	<u>001</u>	Jan 08, 2018
<u>AB</u>		<u>25MG</u>	<u>A209284</u>	<u>002</u>	Jan 08, 2018
<u>AB</u>	HETERO LABS LTD V	<u>12.5MG</u>	<u>A204574</u>	<u>001</u>	Feb 03, 2016
<u>AB</u>		<u>25MG</u>	<u>A204574</u>	<u>002</u>	Feb 03, 2016
<u>AB</u>	MYLAN	<u>12.5MG</u>	<u>A207682</u>	<u>001</u>	Jan 31, 2017
<u>AB</u>		<u>25MG</u>	<u>A207682</u>	<u>002</u>	Jan 31, 2017

XENAZINE

<u>AB</u>	+	BAUSCH	<u>12.5MG</u>	<u>N021894</u>	<u>001</u>	Aug 15, 2008
<u>AB</u>	+!		<u>25MG</u>	<u>N021894</u>	<u>002</u>	Aug 15, 2008

TETRACAINE HYDROCHLORIDE

SOLUTION; OPHTHALMIC

TETRACAINE HYDROCHLORIDE

<u>AB1</u>	+!	BAUSCH LOMB IRELAND	<u>0.5%</u>	<u>N210821</u>	<u>001</u>	Mar 12, 2019
<u>AB1</u>		SOMERSET THERAPS	<u>0.5%</u>	<u>A217227</u>	<u>001</u>	Dec 20, 2024
		LLC				
	+!	ALCON LABS	0.5%	N208135	001	Feb 29, 2016

PRESCRIPTION DRUG PRODUCT LIST

TETRACYCLINE HYDROCHLORIDE

CAPSULE;ORAL

ACHROMYCIN V

AB	+	AVET	250MG	N050278	003	
AB	+		500MG	N050278	001	

TETRACYCLINE HYDROCHLORIDE

AB		AMNEAL PHARMS NY	250MG	A210674	001	Sep 18, 2018
AB	!		500MG	A210674	002	Sep 18, 2018
AB		BRECKENRIDGE	250MG	A210662	001	Nov 07, 2018
AB			500MG	A210662	002	Nov 07, 2018
AB		CHARTWELL TETRA	250MG	A062752	001	Aug 12, 1988
AB			500MG	A062752	002	Aug 12, 1988
AB		ESJAY PHARMA	250MG	A212635	001	Mar 03, 2020
AB			500MG	A212635	002	Mar 03, 2020
AB		WATSON LABS	250MG	A061837	001	
AB			500MG	A061837	002	

TABLET;ORAL

SUMYCIN

		STRIDES PHARMA	250MG	A061147	001	
	!		500MG	A061147	004	

TETRAHYDROZOLINE HYDROCHLORIDE

SOLUTION;NASAL

TYZINE

!	FOUGERA PHARMS	0.05%	A086576	002	
		0.1%	A086576	001	

SPRAY;NASAL

TYZINE

!	FOUGERA PHARMS	0.1%	A086576	003	
---	----------------	------	---------	-----	--

THALIDOMIDE

CAPSULE;ORAL

THALOMID

+	BRISTOL-MYERS	50MG	N020785	001	Jul 16, 1998
+		100MG	N020785	002	Jan 17, 2003
+		150MG	N020785	004	Jan 10, 2007
+	!	200MG	N020785	003	Jan 17, 2003

THALLOUS CHLORIDE TL-201

INJECTABLE;INJECTION

THALLOUS CHLORIDE TL 201

+	CURIUM	1mCi/ML	N018150	001	
---	--------	---------	---------	-----	--

THEOPHYLLINE

CAPSULE, EXTENDED RELEASE;ORAL

THEO-24

	ENDO OPERATIONS	100MG	A087943	002	Aug 22, 1983
+		200MG	A087943	001	Aug 22, 1983
		300MG	A087943	003	Aug 22, 1983
!		400MG	A087943	004	Feb 28, 1992

SOLUTION;ORAL

THEOPHYLLINE

AA	!	CHARTWELL MOLECULAR	80MG/15ML	A091156	001	Apr 13, 2011
AA		TRIS PHARMA INC	80MG/15ML	A091586	001	Jun 15, 2012

SOLUTION, ELIXIR;ORAL

ELIXOPHYLLIN

AA	+	!	NOSTRUM LABS INC	80MG/15ML	A085186	001
-----------	---	---	------------------	------------------	----------------	------------

THEOPHYLLINE

AA		PHARM ASSOC	80MG/15ML	A206344	001	Dec 16, 2016
-----------	--	-------------	------------------	----------------	------------	--------------

TABLET, EXTENDED RELEASE;ORAL

THEOPHYLLINE

AB		ALEMBIC	300MG	A090430	001	Oct 27, 2010
AB	!		450MG	A090430	002	Oct 27, 2010
AB		AMNEAL	300MG	A216276	001	Mar 20, 2023
AB			450MG	A216276	002	Mar 20, 2023
AB		ANNORA PHARMA	300MG	A217422	001	Sep 08, 2023
AB			450MG	A217422	002	Sep 08, 2023
AB		BIONPHARMA	300MG	A216300	001	Apr 26, 2023
AB			450MG	A216300	002	Apr 26, 2023
AB		GLENMARK PHARMS LTD	300MG	A212184	001	Jun 03, 2021
AB			400MG	A090355	001	Jul 13, 2010
AB			450MG	A212184	002	Jun 03, 2021
AB			600MG	A090355	002	Jul 13, 2010
AB		HARMAN FINOCHEM	300MG	A214806	001	Oct 24, 2023

PRESCRIPTION DRUG PRODUCT LIST

THEOPHYLLINE

TABLET, EXTENDED RELEASE;ORAL

THEOPHYLLINE

<u>AB</u>		<u>450MG</u>	<u>A214806 002</u>	Oct 24, 2023
<u>AB</u>	HERITAGE PHARMA	<u>300MG</u>	<u>A089763 001</u>	Apr 30, 1990
	AVET			
<u>AB</u>		<u>450MG</u>	<u>A081236 001</u>	Nov 09, 1992
<u>AB</u>	NOSTRUM LABS INC	<u>400MG</u>	<u>A040560 003</u>	Apr 21, 2006
<u>AB</u>	!	<u>600MG</u>	<u>A040560 002</u>	Apr 21, 2006
<u>AB</u>	NOSTRUM PHARMS LLC	<u>300MG</u>	<u>A087400 002</u>	Jan 11, 1983
<u>AB</u>		<u>450MG</u>	<u>A087400 005</u>	Aug 09, 2023
<u>AB</u>	RHODES PHARMS	<u>300MG</u>	<u>A214113 001</u>	May 11, 2023
<u>AB</u>	+	<u>400MG</u>	<u>A040086 002</u>	Sep 01, 1982
<u>AB</u>	STRIDES PHARMA	<u>300MG</u>	<u>A215312 001</u>	Sep 05, 2024
<u>AB</u>		<u>450MG</u>	<u>A215312 002</u>	Sep 05, 2024
<u>AB</u>	TEVA PHARMS INC	<u>300MG</u>	<u>A216961 001</u>	Oct 12, 2022
<u>AB</u>		<u>450MG</u>	<u>A216961 002</u>	Oct 12, 2022
<u>AB</u>	VELZEN PHARMA PVT	<u>300MG</u>	<u>A214950 001</u>	Jan 30, 2023
<u>AB</u>		<u>450MG</u>	<u>A214950 002</u>	Jan 30, 2023
<u>AB</u>	ZYDUS LIFESCIENCES	<u>300MG</u>	<u>A218063 001</u>	May 21, 2024
<u>AB</u>		<u>450MG</u>	<u>A218063 002</u>	May 21, 2024
	NOSTRUM PHARMS LLC	100MG	A087400 003	Feb 21, 1985
		200MG	A087400 004	Feb 21, 1985

THIAMINE HYDROCHLORIDE

INJECTABLE; INJECTION

THIAMINE HYDROCHLORIDE

<u>AP</u>	ASPIRO	<u>100MG/ML</u>	<u>A218471 001</u>	Nov 06, 2024
<u>AP</u>	CAPLIN	<u>100MG/ML</u>	<u>A215692 001</u>	Mar 06, 2023
<u>AP</u>	DR REDDYS	<u>100MG/ML</u>	<u>A080571 001</u>	
<u>AP</u>	EUGIA PHARMA	<u>100MG/ML</u>	<u>A208703 001</u>	Apr 20, 2022
<u>AP</u>	+!	<u>100MG/ML</u>	<u>A080556 001</u>	
<u>AP</u>	MYLAN INSTITUTIONAL	<u>100MG/ML</u>	<u>A091623 001</u>	Jun 25, 2012
<u>AP</u>	SAGENT PHARMS INC	<u>100MG/ML</u>	<u>A206106 001</u>	Dec 01, 2017
<u>AP</u>	STERISCIENCE	<u>100MG/ML</u>	<u>A217181 001</u>	Oct 29, 2024
	SPECLTS			
<u>AP</u>	WEST-WARD PHARMS	<u>100MG/ML</u>	<u>A080575 001</u>	
	INT			
	!	DR REDDYS	A080571 002	200MG/ML

THIOGUANINE

TABLET; ORAL

THIOGUANINE

+! WAYLIS THERAP 40MG N012429 001

THIORIDAZINE HYDROCHLORIDE

TABLET; ORAL

THIORIDAZINE HYDROCHLORIDE

MYLAN	10MG	A088004 002	Mar 15, 1983
	25MG	A088004 003	Mar 15, 1983
	50MG	A088004 004	Mar 15, 1983
!	100MG	A088004 001	Nov 18, 1983

THIOTEPA

INJECTABLE; INJECTION

THIOTEPA

<u>AP</u>	BELOTECA	<u>15MG/VIAL</u>	<u>A211831 001</u>	Dec 08, 2021
<u>AP</u>	DR REDDYS	<u>15MG/VIAL</u>	<u>A210337 001</u>	May 04, 2018
<u>AP</u>	HENGRUI PHARMA	<u>15MG/VIAL</u>	<u>A209150 001</u>	May 04, 2018
<u>AP</u>	PENN LIFE	<u>15MG/VIAL</u>	<u>A208242 001</u>	Jan 10, 2020
<u>AP</u>	!	WEST-WARD PHARMS	<u>A075547 001</u>	Apr 02, 2001
	INT			

POWDER; INTRACAVITARY, INTRAVENOUS, INTRAVESICAL

TEPADINA

<u>AP</u>	+!	ADIENNE SA	<u>15MG/VIAL</u>	<u>N208264 001</u>	Jan 26, 2017
<u>AP</u>	+!		<u>100MG/VIAL</u>	<u>N208264 002</u>	Jan 26, 2017

THIOTEPA

<u>AP</u>	BELOTECA	<u>15MG/VIAL</u>	<u>A217559 001</u>	Aug 20, 2024
<u>AP</u>	GLAND PHARMA LTD	<u>15MG/VIAL</u>	<u>A214222 001</u>	Mar 08, 2021
<u>AP</u>		<u>100MG/VIAL</u>	<u>A214222 002</u>	Jan 03, 2022
<u>AP</u>	HIKMA	<u>100MG/VIAL</u>	<u>A211755 001</u>	Sep 05, 2023
<u>AP</u>	MEITHEAL	<u>15MG/VIAL</u>	<u>A216037 001</u>	Dec 26, 2023
<u>AP</u>		<u>100MG/VIAL</u>	<u>A216037 002</u>	Dec 26, 2023
<u>AP</u>	MSN	<u>15MG/VIAL</u>	<u>A213049 001</u>	Mar 04, 2020
<u>AP</u>		<u>100MG/VIAL</u>	<u>A213049 002</u>	Mar 04, 2020

PRESCRIPTION DRUG PRODUCT LIST

THIOTEPA

SOLUTION; INTRAVENOUS

TEPYLUTE

+! SHORLA

15MG/1.5ML (10MG/ML)

N216984 001 Jun 25, 2024

THIOTHIXENE

CAPSULE; ORAL

THIOTHIXENEAB AMNEAL1MGA215456 001 Feb 28, 2022AB2MGA215456 002 Feb 28, 2022AB5MGA215456 003 Feb 28, 2022AB10MGA215456 004 Feb 28, 2022AB NOVITIUM PHARMA1MGA211642 001 Apr 05, 2019AB2MGA211642 002 Apr 05, 2019AB !5MGA211642 003 Apr 05, 2019AB10MGA211642 004 Apr 05, 2019AB RISING1MGA071093 002 Jun 23, 1987AB2MGA071093 003 Jun 23, 1987AB5MGA071093 004 Jun 23, 1987AB10MGA071093 001 Jun 23, 1987TIAGABINE HYDROCHLORIDE

TABLET; ORAL

GABITRILAB + CEPHALON2MGN020646 005 Apr 16, 1999AB +!4MGN020646 001 Sep 30, 1997AB +12MGN020646 002 Sep 30, 1997AB +16MGN020646 003 Sep 30, 1997TIAGABINE HYDROCHLORIDEAB MSN2MGA214816 001 Nov 16, 2021AB4MGA214816 002 Nov 16, 2021AB12MGA214816 003 Nov 16, 2021AB16MGA214816 004 Nov 16, 2021AB SUN PHARM INDS2MGA077555 001 Nov 04, 2011AB4MGA077555 002 Nov 04, 2011TICAGRELOR

TABLET; ORAL

BRILINTAAB +! ASTRAZENECA90MGN022433 001 Jul 20, 2011TICAGRELORAB ALKEM LABS LTD90MGA208567 001 Apr 21, 2023AB DR REDDYS90MGA208541 001 Oct 29, 2024AB HISUN PHARM90MGA208575 001 Jan 23, 2019

HANGZHOU

AB PRINSTON INC90MGA208599 001 Aug 16, 2024

BRILINTA

+ ASTRAZENECA

60MG

N022433 002 Sep 03, 2015

TICLOPIDINE HYDROCHLORIDE

TABLET; ORAL

TICLOPIDINE HYDROCHLORIDE

! APOTEX

250MG

A075089 001 Jul 01, 1999

TIGECYCLINE

POWDER; INTRAVENOUS

TIGECYCLINEAP AMNEAL50MG/VIALN211158 001 Aug 02, 2018AP APOTEX50MG/VIALA204439 001 Dec 21, 2018AP EUGIA PHARMA50MG/VIALA206335 001 Jun 11, 2019AP +! FRESENIUS KABI USA50MG/VIALN205645 001 Dec 01, 2016AP MEITHEAL50MG/VIALA214020 001 May 13, 2021AP SANDOZ50MG/VIALA091620 001 May 27, 2015TYGACILAP +! PF PRISM CV50MG/VIALN021821 001 Jun 15, 2005TIMOLOL

SOLUTION/DROPS; OPHTHALMIC

BETIMOLAT +! THEA PHARMA0.5%N020439 002 Mar 31, 1995TIMOLOLAT SOMERSET0.5%A216654 001 Oct 10, 2024

BETIMOL

+! THEA PHARMA

0.25%

N020439 001 Mar 31, 1995

PRESCRIPTION DRUG PRODUCT LIST

TIMOLOL MALEATE

SOLUTION, GEL FORMING/DROPS;OPHTHALMIC

TIMOLOL MALEATE

<u>AB</u>	!	ALEMBIC	<u>EQ 0.25% BASE</u>	<u>A212942 001</u>	Oct 22, 2020
<u>AB</u>	!		<u>EQ 0.5% BASE</u>	<u>A212942 002</u>	Oct 22, 2020
<u>AB</u>		DR REDDYS	<u>EQ 0.25% BASE</u>	<u>A215733 001</u>	Sep 22, 2022
<u>AB</u>			<u>EQ 0.5% BASE</u>	<u>A215733 002</u>	Sep 22, 2022
<u>AB</u>		EUGIA PHARMA	<u>EQ 0.5% BASE</u>	<u>A213540 001</u>	Mar 23, 2023
<u>AB</u>		GLAND PHARMA LTD	<u>EQ 0.25% BASE</u>	<u>A214645 001</u>	Jun 24, 2022
<u>AB</u>			<u>EQ 0.5% BASE</u>	<u>A214645 002</u>	Feb 14, 2023
<u>AB</u>	+	SANDOZ	<u>EQ 0.25% BASE</u>	<u>N020963 001</u>	Oct 21, 1998
<u>AB</u>	+		<u>EQ 0.5% BASE</u>	<u>N020963 002</u>	Oct 21, 1998
<u>AB</u>		SOMERSET THERAPS LLC	<u>EQ 0.25% BASE</u>	<u>A210640 001</u>	Sep 16, 2024
<u>AB</u>			<u>EQ 0.5% BASE</u>	<u>A210640 002</u>	Sep 16, 2024
<u>TIMOPTIC-XE</u>					
<u>AB</u>	+	BAUSCH AND LOMB INC	<u>EQ 0.25% BASE</u>	<u>N020330 001</u>	Nov 04, 1993
<u>AB</u>	+		<u>EQ 0.5% BASE</u>	<u>N020330 002</u>	Nov 04, 1993

SOLUTION/DROPS;OPHTHALMIC

TIMOLOL MALEATE

<u>AT1</u>		CAPLIN	<u>EQ 0.5% BASE</u>	<u>A218460 001</u>	Aug 07, 2024
<u>AT1</u>		FDC LTD	<u>EQ 0.25% BASE</u>	<u>A077259 001</u>	Apr 30, 2008
<u>AT1</u>			<u>EQ 0.5% BASE</u>	<u>A077259 002</u>	Apr 30, 2008
<u>AT1</u>		MANKIND PHARMA	<u>EQ 0.25% BASE</u>	<u>A078771 001</u>	Sep 28, 2009
<u>AT1</u>			<u>EQ 0.5% BASE</u>	<u>A078771 002</u>	Sep 28, 2009
<u>AT1</u>		MICRO LABS	<u>EQ 0.25% BASE</u>	<u>A217343 001</u>	May 26, 2023
<u>AT1</u>			<u>EQ 0.5% BASE</u>	<u>A217343 002</u>	May 26, 2023
<u>AT1</u>		PACIFIC PHARMA	<u>EQ 0.25% BASE</u>	<u>A074746 001</u>	Mar 25, 1997
<u>AT1</u>			<u>EQ 0.5% BASE</u>	<u>A074747 001</u>	Mar 25, 1997
<u>AT1</u>	!	SANDOZ	<u>EQ 0.25% BASE</u>	<u>A074261 001</u>	Apr 28, 1995
<u>AT1</u>	!		<u>EQ 0.5% BASE</u>	<u>A074262 001</u>	Apr 28, 1995
<u>AT1</u>		SOMERSET	<u>EQ 0.25% BASE</u>	<u>A217764 002</u>	Jul 03, 2024
<u>AT1</u>			<u>EQ 0.5% BASE</u>	<u>A217764 001</u>	Jul 03, 2024

TIMOPTIC

<u>AT1</u>	+	BAUSCH AND LOMB INC	<u>EQ 0.25% BASE</u>	<u>N018086 001</u>	
<u>AT1</u>	+		<u>EQ 0.5% BASE</u>	<u>N018086 002</u>	

ISTALOL

<u>AT2</u>	+	BAUSCH AND LOMB	<u>EQ 0.5% BASE</u>	<u>N021516 001</u>	Jun 04, 2004
------------	---	-----------------	---------------------	--------------------	--------------

TIMOLOL MALEATE

<u>AT2</u>		APOTEX	<u>EQ 0.5% BASE</u>	<u>A204936 001</u>	Apr 17, 2015
<u>AT2</u>		SOMERSET THERAPS LLC	<u>EQ 0.5% BASE</u>	<u>A216653 001</u>	Jul 03, 2024
<u>AT3</u>		AMRING PHARMS	<u>EQ 0.25% BASE</u>	<u>A212592 001</u>	Dec 13, 2021
<u>AT3</u>			<u>EQ 0.5% BASE</u>	<u>A212592 002</u>	Dec 13, 2021
<u>AT3</u>		EPIC PHARMA LLC	<u>EQ 0.5% BASE</u>	<u>A212291 001</u>	Sep 11, 2020
<u>AT3</u>		INGENUS PHARMS LLC	<u>EQ 0.25% BASE</u>	<u>A216533 001</u>	Sep 13, 2022
<u>AT3</u>			<u>EQ 0.5% BASE</u>	<u>A216533 002</u>	Sep 13, 2022
<u>AT3</u>		MICRO LABS	<u>EQ 0.5% BASE</u>	<u>A216596 001</u>	Sep 07, 2022
<u>AT3</u>		SENTISS PHARMA	<u>EQ 0.5% BASE</u>	<u>A217195 001</u>	May 08, 2023

TIMOPTIC IN OCULOSE

<u>AT3</u>	+	BAUSCH AND LOMB INC	<u>EQ 0.25% BASE</u>	<u>N019463 001</u>	Nov 05, 1986
<u>AT3</u>	+		<u>EQ 0.5% BASE</u>	<u>N019463 002</u>	Nov 05, 1986

TABLET;ORAL

TIMOLOL MALEATE

<u>AB</u>		MYLAN	<u>5MG</u>	<u>A072668 002</u>	Jun 08, 1990
<u>AB</u>			<u>10MG</u>	<u>A072668 003</u>	Jun 08, 1990
<u>AB</u>	!		<u>20MG</u>	<u>A072668 001</u>	Jun 08, 1990
<u>AB</u>		RISING	<u>5MG</u>	<u>A207556 001</u>	May 02, 2019
<u>AB</u>			<u>10MG</u>	<u>A207556 002</u>	May 02, 2019
<u>AB</u>			<u>20MG</u>	<u>A207556 003</u>	May 02, 2019

TINIDAZOLE

TABLET;ORAL

TINDAMAX

<u>AB</u>	+	MISSION PHARMA	<u>250MG</u>	<u>N021618 001</u>	May 17, 2004
<u>AB</u>	+		<u>500MG</u>	<u>N021618 002</u>	May 17, 2004

TINIDAZOLE

<u>AB</u>		CHARTWELL RX	<u>250MG</u>	<u>A202044 001</u>	Apr 30, 2012
<u>AB</u>			<u>500MG</u>	<u>A202044 002</u>	Apr 30, 2012
<u>AB</u>		EDENBRIDGE PHARMS	<u>250MG</u>	<u>A203808 001</u>	Aug 04, 2015
<u>AB</u>			<u>500MG</u>	<u>A203808 002</u>	Aug 04, 2015
<u>AB</u>		THINQ PHARM-CRO PVT	<u>250MG</u>	<u>A202489 001</u>	Oct 09, 2013
<u>AB</u>			<u>500MG</u>	<u>A202489 002</u>	Oct 09, 2013

PRESCRIPTION DRUG PRODUCT LIST

TIOPRONIN

TABLET; ORAL

THIOLA

AB	+	MISSION PHARMA	100MG	N019569	001	Aug 11, 1988
-----------	----------	----------------	--------------	----------------	------------	--------------

TIOPRONIN

AB		TEVA PHARMS USA INC	100MG	A214326	001	Apr 26, 2021
-----------	--	---------------------	--------------	----------------	------------	--------------

TABLET, DELAYED RELEASE; ORAL

THIOLA EC

AB	+	MISSION PHARMACAL	100MG	N211843	001	Jun 28, 2019
-----------	----------	-------------------	--------------	----------------	------------	--------------

AB	+		300MG	N211843	002	Jun 28, 2019
-----------	----------	--	--------------	----------------	------------	--------------

TIOPRONIN

AB		AMNEAL	300MG	A216278	001	Aug 15, 2023
-----------	--	--------	--------------	----------------	------------	--------------

AB		ENDO OPERATIONS	100MG	A217219	001	Feb 24, 2023
-----------	--	-----------------	--------------	----------------	------------	--------------

AB			300MG	A217219	002	Feb 24, 2023
-----------	--	--	--------------	----------------	------------	--------------

AB		TEVA PHARMS INC	100MG	A216456	001	Jul 22, 2024
-----------	--	-----------------	--------------	----------------	------------	--------------

AB			300MG	A216456	002	Jul 22, 2024
-----------	--	--	--------------	----------------	------------	--------------

AB		TORRENT	100MG	A216990	001	Jan 30, 2024
-----------	--	---------	--------------	----------------	------------	--------------

AB			300MG	A216990	002	Jan 30, 2024
-----------	--	--	--------------	----------------	------------	--------------

TIOTROPIUM BROMIDE

POWDER; INHALATION

SPIRIVA

AB	+	BOEHRINGER INGELHEIM	EQ 0.018MG BASE/INH	N021395	001	Jan 30, 2004
-----------	----------	-------------------------	----------------------------	----------------	------------	--------------

TIOTROPIUM BROMIDE

AB		LUPIN	EQ 0.018MG BASE/INH	A211287	001	Jun 20, 2023
-----------	--	-------	----------------------------	----------------	------------	--------------

SPRAY, METERED; INHALATION

SPIRIVA RESPIMAT

+	!	BOEHRINGER INGELHEIM	EQ 0.00125MG BASE/INH	N021936	002	Sep 15, 2015
----------	----------	-------------------------	------------------------------	----------------	------------	--------------

+	!		EQ 0.0025MG BASE/INH	N021936	001	Sep 24, 2014
----------	----------	--	-----------------------------	----------------	------------	--------------

TIPIRACIL HYDROCHLORIDE; TRIFLURIDINE

TABLET; ORAL

LONSURF

AB	+	TAIHO ONCOLOGY	EQ 6.14MG BASE;15MG	N207981	001	Sep 22, 2015
-----------	----------	----------------	----------------------------	----------------	------------	--------------

AB	+		EQ 8.19MG BASE;20MG	N207981	002	Sep 22, 2015
-----------	----------	--	----------------------------	----------------	------------	--------------

TIPIRACIL HYDROCHLORIDE AND TRIFLURIDINE

AB		MSN	EQ 6.14MG BASE;15MG	A214024	001	Jan 08, 2025
-----------	--	-----	----------------------------	----------------	------------	--------------

AB			EQ 8.19MG BASE;20MG	A214024	002	Jan 08, 2025
-----------	--	--	----------------------------	----------------	------------	--------------

AB		NATCO	EQ 6.14MG BASE;15MG	A214008	001	Jun 13, 2023
-----------	--	-------	----------------------------	----------------	------------	--------------

AB			EQ 8.19MG BASE;20MG	A214008	002	Jun 13, 2023
-----------	--	--	----------------------------	----------------	------------	--------------

TIPRANAVIR

CAPSULE; ORAL

APTIVUS

+	!	BOEHRINGER INGELHEIM	250MG	N021814	001	Jun 22, 2005
----------	----------	-------------------------	--------------	----------------	------------	--------------

TIRBANIBULIN

OINTMENT; TOPICAL

KLISYRI

+	!	ALMIRALL	1%	N213189	001	Dec 14, 2020
----------	----------	----------	-----------	----------------	------------	--------------

TIROFIBAN HYDROCHLORIDE

SOLUTION; INJECTION

AGGRASTAT

+	!	MEDICURE	EQ 3.75MG BASE/15ML (EQ 0.25MG BASE/ML)	N020912	002	Aug 31, 2016
----------	----------	----------	--	----------------	------------	--------------

SOLUTION; INTRAVENOUS

AGGRASTAT

AP	+	MEDICURE	EQ 5MG BASE/100ML (EQ 0.05MG BASE/ML)	N020913	002	May 17, 2002
-----------	----------	----------	--	----------------	------------	--------------

AP	+		EQ 12.5MG BASE/250ML (EQ 0.05MG BASE/ML)	N020913	003	Apr 20, 2000
-----------	----------	--	---	----------------	------------	--------------

TIROFIBAN HYDROCHLORIDE

AP		EUGIA PHARMA	EQ 5MG BASE/100ML (EQ 0.05MG BASE/ML)	A216379	001	May 01, 2023
-----------	--	--------------	--	----------------	------------	--------------

AP			EQ 12.5MG BASE/250ML (EQ 0.05MG BASE/ML)	A216379	002	May 01, 2023
-----------	--	--	---	----------------	------------	--------------

AP		GLAND PHARMA LTD	EQ 12.5MG BASE/250ML (EQ 0.05MG BASE/ML)	A206888	001	Apr 08, 2021
-----------	--	------------------	---	----------------	------------	--------------

AP		NEXUS	EQ 5MG BASE/100ML (EQ 0.05MG BASE/ML)	A213947	001	Feb 07, 2023
-----------	--	-------	--	----------------	------------	--------------

AP			EQ 12.5MG BASE/250ML (EQ 0.05MG BASE/ML)	A213947	002	Jul 24, 2023
-----------	--	--	---	----------------	------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

TIRZEPATIDE

SOLUTION;SUBCUTANEOUS

MOUNJARO

+	!	ELI LILLY AND CO	2.5MG/0.5ML (2.5MG/0.5ML)	N215866	007	Jul 28, 2023
+	!		5MG/0.5ML (5MG/0.5ML)	N215866	008	Jul 28, 2023
+	!		7.5MG/0.5ML (7.5MG/0.5ML)	N215866	009	Jul 28, 2023
+	!		10MG/0.5ML (10MG/0.5ML)	N215866	010	Jul 28, 2023
+	!		12.5MG/0.5ML (12.5MG/0.5ML)	N215866	011	Jul 28, 2023
+	!		15MG/0.5ML (15MG/0.5ML)	N215866	012	Jul 28, 2023

MOUNJARO (AUTOINJECTOR)

+	!	ELI LILLY AND CO	2.5MG/0.5ML (2.5MG/0.5ML)	N215866	001	May 13, 2022
+	!		5MG/0.5ML (5MG/0.5ML)	N215866	002	May 13, 2022
+	!		7.5MG/0.5ML (7.5MG/0.5ML)	N215866	003	May 13, 2022
+	!		10MG/0.5ML (10MG/0.5ML)	N215866	004	May 13, 2022
+	!		12.5MG/0.5ML (12.5MG/0.5ML)	N215866	005	May 13, 2022
+	!		15MG/0.5ML (15MG/0.5ML)	N215866	006	May 13, 2022

ZEPBOUND

+	!	ELI LILLY AND CO	2.5MG/0.5ML (2.5MG/0.5ML)	N217806	007	Mar 28, 2024
+	!		5MG/0.5ML (5MG/0.5ML)	N217806	008	Mar 28, 2024
+	!		7.5MG/0.5ML (7.5MG/0.5ML)	N217806	009	Mar 28, 2024
+	!		10MG/0.5ML (10MG/0.5ML)	N217806	010	Mar 28, 2024
+	!		12.5MG/0.5ML (12.5MG/0.5ML)	N217806	011	Mar 28, 2024
+	!		15MG/0.5ML (15MG/0.5ML)	N217806	012	Mar 28, 2024

ZEPBOUND (AUTOINJECTOR)

+	!	ELI LILLY AND CO	2.5MG/0.5ML (2.5MG/0.5ML)	N217806	001	Nov 08, 2023
+	!		5MG/0.5ML (5MG/0.5ML)	N217806	002	Nov 08, 2023
+	!		7.5MG/0.5ML (7.5MG/0.5ML)	N217806	003	Nov 08, 2023
+	!		10MG/0.5ML (10MG/0.5ML)	N217806	004	Nov 08, 2023
+	!		12.5MG/0.5ML (12.5MG/0.5ML)	N217806	005	Nov 08, 2023
+	!		15MG/0.5ML (15MG/0.5ML)	N217806	006	Nov 08, 2023

TIVOZANIB HYDROCHLORIDE

CAPSULE;ORAL

FOTIVDA

+		AVEO PHARMS	EQ 0.89MG BASE	N212904	001	Mar 10, 2021
+	!		EQ 1.34MG BASE	N212904	002	Mar 10, 2021

TIZANIDINE HYDROCHLORIDE

CAPSULE;ORAL

TIZANIDINE HYDROCHLORIDE

<u>AB</u>		ALEMbic	<u>EQ 2MG BASE</u>	<u>A213223</u>	<u>001</u>	Jan 13, 2020
<u>AB</u>			<u>EQ 4MG BASE</u>	<u>A213223</u>	<u>002</u>	Jan 13, 2020
<u>AB</u>			<u>EQ 6MG BASE</u>	<u>A213223</u>	<u>003</u>	Jan 13, 2020
<u>AB</u>		ALKEM LABS LTD	<u>EQ 2MG BASE</u>	<u>A212196</u>	<u>001</u>	Mar 27, 2019
<u>AB</u>			<u>EQ 4MG BASE</u>	<u>A212196</u>	<u>002</u>	Mar 27, 2019
<u>AB</u>			<u>EQ 6MG BASE</u>	<u>A212196</u>	<u>003</u>	Mar 27, 2019
<u>AB</u>		APOTEX INC	<u>EQ 2MG BASE</u>	<u>A078868</u>	<u>001</u>	Feb 03, 2012
<u>AB</u>			<u>EQ 4MG BASE</u>	<u>A078868</u>	<u>002</u>	Feb 03, 2012
<u>AB</u>			<u>EQ 6MG BASE</u>	<u>A078868</u>	<u>003</u>	Feb 03, 2012
<u>AB</u>		AUROBINDO PHARMA LTD	<u>EQ 2MG BASE</u>	<u>A213544</u>	<u>001</u>	Mar 24, 2020
<u>AB</u>			<u>EQ 4MG BASE</u>	<u>A213544</u>	<u>002</u>	Mar 24, 2020
<u>AB</u>			<u>EQ 6MG BASE</u>	<u>A213544</u>	<u>003</u>	Mar 24, 2020
<u>AB</u>		CADILA PHARMS LTD	<u>EQ 2MG BASE</u>	<u>A210021</u>	<u>001</u>	Mar 26, 2019
<u>AB</u>			<u>EQ 4MG BASE</u>	<u>A210021</u>	<u>002</u>	Mar 26, 2019
<u>AB</u>			<u>EQ 6MG BASE</u>	<u>A210021</u>	<u>003</u>	Mar 26, 2019
<u>AB</u>		JUBILANT GENERICS	<u>EQ 2MG BASE</u>	<u>A209605</u>	<u>001</u>	Aug 04, 2017
<u>AB</u>			<u>EQ 4MG BASE</u>	<u>A209605</u>	<u>002</u>	Aug 04, 2017
<u>AB</u>			<u>EQ 6MG BASE</u>	<u>A209605</u>	<u>003</u>	Aug 04, 2017
<u>AB</u>		NOVAST LABS	<u>EQ 2MG BASE</u>	<u>A210267</u>	<u>001</u>	Mar 12, 2019
<u>AB</u>			<u>EQ 4MG BASE</u>	<u>A210267</u>	<u>002</u>	Mar 12, 2019
<u>AB</u>			<u>EQ 6MG BASE</u>	<u>A210267</u>	<u>003</u>	Mar 12, 2019
<u>AB</u>		RUBICON	<u>EQ 2MG BASE</u>	<u>A213798</u>	<u>001</u>	May 27, 2020
<u>AB</u>			<u>EQ 4MG BASE</u>	<u>A213798</u>	<u>002</u>	May 27, 2020
<u>AB</u>			<u>EQ 6MG BASE</u>	<u>A213798</u>	<u>003</u>	May 27, 2020
<u>AB</u>		SOMERSET THERAPS LLC	<u>EQ 4MG BASE</u>	<u>A213365</u>	<u>001</u>	May 13, 2024
<u>AB</u>			<u>EQ 6MG BASE</u>	<u>A213365</u>	<u>002</u>	May 13, 2024
<u>AB</u>		ZYDUS PHARMS	<u>EQ 2MG BASE</u>	<u>A208622</u>	<u>001</u>	Mar 03, 2017
<u>AB</u>			<u>EQ 4MG BASE</u>	<u>A208622</u>	<u>002</u>	Mar 03, 2017
<u>AB</u>			<u>EQ 6MG BASE</u>	<u>A208622</u>	<u>003</u>	Mar 03, 2017
<u>AB</u>		<u>ZANAFLEX</u>				
<u>AB</u>	+	LEGACY PHARMA USA	<u>EQ 2MG BASE</u>	<u>N021447</u>	<u>001</u>	Aug 29, 2002
<u>AB</u>	+		<u>EQ 4MG BASE</u>	<u>N021447</u>	<u>002</u>	Aug 29, 2002

PRESCRIPTION DRUG PRODUCT LIST

TIZANIDINE HYDROCHLORIDE

CAPSULE; ORAL

ZANAFLEX**AB** **+**! **EQ 6MG BASE** **N021447 003** Aug 29, 2002

SOLUTION; ORAL

ONTRALFY

+! FIDELITY BIOPHARMA **EQ 2MG BASE/5ML** **N216190 001** Dec 12, 2024

TABLET; ORAL

TIZANIDINE HYDROCHLORIDE**AB** ALKEM LABS LTD **EQ 2MG BASE** **A211798 001** Jan 25, 2019**AB** **EQ 4MG BASE** **A211798 002** Jan 25, 2019**AB** APOTEX **EQ 2MG BASE** **A076533 001** Jan 16, 2004**AB** **EQ 4MG BASE** **A076533 002** Jan 16, 2004**AB** CADILA **EQ 2MG BASE** **A208187 001** Mar 09, 2018**AB** **EQ 4MG BASE** **A208187 002** Mar 09, 2018**AB** DR REDDYS LABS INC **EQ 2MG BASE** **A076286 001** Jul 03, 2002**AB** **EQ 4MG BASE** **A076286 002** Jul 03, 2002**AB** EPIC PHARMA LLC **EQ 2MG BASE** **A076347 001** Oct 11, 2002**AB** **EQ 4MG BASE** **A076347 002** Oct 11, 2002**AB** GRAVITI PHARMS **EQ 4MG BASE** **A218920 002** Aug 07, 2024**AB** **EQ 2MG BASE** **A218920 001** Aug 07, 2024**AB** OXFORD PHARMS **EQ 2MG BASE** **A076281 001** Oct 20, 2003**AB** **EQ 4MG BASE** **A076281 002** Oct 20, 2003**AB** SUN PHARM INDS INC **EQ 2MG BASE** **A076416 001** Sep 29, 2003**AB** **EQ 4MG BASE** **A076416 002** Sep 29, 2003**AB** UNICHEM LABS LTD **EQ 2MG BASE** **A091283 001** Nov 28, 2012**AB** **EQ 4MG BASE** **A091283 002** Nov 28, 2012ZANAFLEX**AB** **+**! LEGACY PHARMA USA **EQ 4MG BASE** **N020397 001** Nov 27, 1996TOBRAMYCIN

OINTMENT; OPHTHALMIC

TOBREX

+! NOVARTIS **0.3%** **N050555 001**

POWDER; INHALATION

TOBI PODHALER

+! MYLAN SPECIALITY LP **28MG** **N201688 001** Mar 22, 2013

SOLUTION; INHALATION

BETHKIS**AN** **+**! CHIESI **300MG/4ML** **N201820 001** Oct 12, 2012KITABIS PAK**AN** PULMOFLOW INC **300MG/5ML** **N205433 001** Dec 02, 2014TOBI**AN** **+**! MYLAN SPECIALITY LP **300MG/5ML** **N050753 001** Dec 22, 1997TOBRAMYCIN**AN** ALKEM LABS LTD **300MG/5ML** **A212848 001** Apr 01, 2021**AN** AMNEAL PHARMS **300MG/5ML** **A205501 001** Jul 13, 2015**AN** DR REDDYS LABS SA **300MG/5ML** **A207080 001** Jul 09, 2018**AN** HIKMA **300MG/5ML** **A201422 001** May 28, 2014**AN** LUPIN **300MG/5ML** **A208964 001** Mar 22, 2017**AN** MANKIND PHARMA **300MG/5ML** **A214478 001** Jun 30, 2023**AN** **300MG/4ML** **A216725 001** Sep 22, 2022**AN** MICRO LABS **300MG/5ML** **A217344 001** Jul 27, 2023**AN** SUN PHARM **300MG/5ML** **A207136 001** Dec 26, 2019**AN** TEVA PHARMS USA **300MG/5ML** **A091589 001** Oct 10, 2013**AN** **300MG/4ML** **A210915 001** Jun 26, 2019

SOLUTION/DROPS; OPHTHALMIC

TOBRAMYCIN**AT** ALEMBIC **0.3%** **A211847 001** Apr 19, 2019**AT** BAUSCH AND LOMB **0.3%** **A064052 001** Nov 29, 1993**AT** CHARTWELL RX **0.3%** **A065026 001** Sep 11, 2001**AT** GLAND PHARMA LTD **0.3%** **A212628 001** Mar 09, 2021**AT** ! SOMERSET THERAPS **0.3%** **A207444 001** Jun 28, 2017

LLC

TOBREX**AT** SANDOZ **0.3%** **A062535 001** Dec 13, 1984TOBRAMYCIN SULFATE

INJECTABLE; INJECTION

TOBRAMYCIN SULFATE**AP** BAXTER HLTHCARE **EQ 40MG BASE/ML** **A206965 001** Jul 01, 2016

CORP

AP EUGIA PHARMA **EQ 1.2GM BASE/VIAL** **A217519 001** Jun 21, 2023**AP** ! FRESENIUS KABI USA **EQ 10MG BASE/ML** **A065122 001** Nov 29, 2002

PRESCRIPTION DRUG PRODUCT LIST

TOBRAMYCIN SULFATE

INJECTABLE; INJECTION

TOBRAMYCIN SULFATE

AP	!		<u>EQ 40MG BASE/ML</u>	<u>A065122 002</u>	Nov 29, 2002
AP			<u>EQ 1.2GM BASE/VIAL</u>	<u>N050789 001</u>	Jul 13, 2004
AP		GLAND PHARMA LTD	<u>EQ 10MG BASE/ML</u>	<u>A209621 001</u>	Feb 11, 2021
AP			<u>EQ 40MG BASE/ML</u>	<u>A209621 002</u>	Feb 11, 2021
AP			<u>EQ 1.2GM BASE/VIAL</u>	<u>A211189 001</u>	May 18, 2021
AP		HIKMA	<u>EQ 40MG BASE/ML</u>	<u>A063117 001</u>	Apr 26, 1991
AP		HOSPIRA	<u>EQ 40MG BASE/ML</u>	<u>A063111 001</u>	Apr 30, 1991
AP		MYLAN LABS LTD	<u>EQ 40MG BASE/ML</u>	<u>A065407 001</u>	Mar 11, 2008
AP		SLATE RUN PHARMA	<u>EQ 1.2GM BASE/VIAL</u>	<u>A217029 001</u>	Feb 01, 2023
AP		XELLIA PHARMS APS	<u>EQ 1.2GM BASE/VIAL</u>	<u>A205685 001</u>	Sep 16, 2014
AP	!	XGEN PHARMS	<u>EQ 1.2GM BASE/VIAL</u>	<u>A065013 001</u>	Aug 17, 2001
<u>TOBRAMYCIN SULFATE (PHARMACY BULK)</u>					
AP	!	FRESENIUS KABI USA	<u>EQ 40MG BASE/ML</u>	<u>A065120 001</u>	Nov 29, 2002
AP		GLAND PHARMA LTD	<u>EQ 40MG BASE/ML</u>	<u>A209346 001</u>	Feb 09, 2021

TOFACITINIB CITRATE

SOLUTION; ORAL

XELJANZ

+! PFIZER

EQ 1MG BASE/ML

N213082 001 Sep 25, 2020

TABLET; ORAL

XELJANZ

+ PF PRISM CV

EQ 5MG BASE

N203214 001 Nov 06, 2012

+!

EQ 10MG BASE

N203214 002 May 30, 2018

TABLET, EXTENDED RELEASE; ORAL

TOFACITINIB CITRATE

AB		DEXCEL	<u>EQ 11MG BASE</u>	<u>A218668 001</u>	Dec 16, 2024
<u>XELJANZ XR</u>					
AB	+	PFIZER	<u>EQ 11MG BASE</u>	<u>N208246 001</u>	Feb 23, 2016
		!	EQ 22MG BASE	<u>N208246 002</u>	Dec 12, 2019

TOFERSEN

SOLUTION; INTRATHECAL

QALSODY

+! BIOGEN MA

100MG/15ML (6.7MG/ML)

N215887 001 Apr 25, 2023

TOLCAPONE

TABLET; ORAL

TASMAR

AB	+	BAUSCH	<u>100MG</u>	<u>N020697 001</u>	Jan 29, 1998
-----------	---	--------	--------------	--------------------	--------------

TOLCAPONE

AB		NOVAST LABS	<u>100MG</u>	<u>A208937 001</u>	Aug 07, 2018
-----------	--	-------------	--------------	--------------------	--------------

TOLMETIN SODIUM

CAPSULE; ORAL

TOLMETIN SODIUM

! RISING

EQ 400MG BASE

A073393 001 May 27, 1993

TABLET; ORAL

TOLMETIN SODIUM

! RISING

EQ 600MG BASE

A074473 001 Aug 30, 1994

TOLTERODINE TARTRATE

CAPSULE, EXTENDED RELEASE; ORAL

DETROL LA

AB	+	UPJOHN	<u>2MG</u>	<u>N021228 001</u>	Dec 22, 2000
AB	+	!	<u>4MG</u>	<u>N021228 002</u>	Dec 22, 2000

TOLTERODINE TARTRATE

AB		AJANTA PHARMA LTD	<u>2MG</u>	<u>A213397 001</u>	May 19, 2020
AB			<u>4MG</u>	<u>A213397 002</u>	May 19, 2020
AB		HETERO LABS LTD III	<u>2MG</u>	<u>A206419 001</u>	Dec 12, 2017
AB			<u>4MG</u>	<u>A206419 002</u>	Dec 12, 2017
AB		INVENTIA HLTHCARE	<u>2MG</u>	<u>A204562 001</u>	Aug 19, 2019
AB			<u>4MG</u>	<u>A204562 002</u>	Aug 19, 2019
AB		TEVA PHARMS USA	<u>2MG</u>	<u>A079141 001</u>	Nov 22, 2016
AB			<u>4MG</u>	<u>A079141 002</u>	Nov 22, 2016
AB		TORRENT	<u>2MG</u>	<u>A203016 001</u>	Aug 11, 2015
AB			<u>4MG</u>	<u>A203016 002</u>	Aug 11, 2015
AB		UNICHEM	<u>2MG</u>	<u>A216917 001</u>	Aug 27, 2024
AB			<u>4MG</u>	<u>A216917 002</u>	Aug 27, 2024
AB		UTOPIC PHARMS	<u>2MG</u>	<u>A213858 001</u>	Feb 02, 2021
AB			<u>4MG</u>	<u>A213858 002</u>	Feb 02, 2021

PRESCRIPTION DRUG PRODUCT LIST

TOLTERODINE TARTRATE

TABLET;ORAL

DETROL

<u>AB</u>	+	UPJOHN	<u>1MG</u>	<u>N020771</u>	<u>001</u>	Mar 25, 1998
<u>AB</u>	+	!	<u>2MG</u>	<u>N020771</u>	<u>002</u>	Mar 25, 1998

TOLTERODINE TARTRATE

<u>AB</u>		ELYSIUM	<u>1MG</u>	<u>A210775</u>	<u>001</u>	Dec 30, 2019
<u>AB</u>			<u>2MG</u>	<u>A210775</u>	<u>002</u>	Dec 30, 2019
<u>AB</u>		HETERO LABS LTD V	<u>1MG</u>	<u>A204397</u>	<u>001</u>	Aug 02, 2021
<u>AB</u>			<u>2MG</u>	<u>A204397</u>	<u>002</u>	Aug 02, 2021
<u>AB</u>		IVAX SUB TEVA PHARMS	<u>1MG</u>	<u>A077006</u>	<u>001</u>	Feb 23, 2015
<u>AB</u>			<u>2MG</u>	<u>A077006</u>	<u>002</u>	Feb 23, 2015
<u>AB</u>		MACLEODS PHARMS LTD	<u>1MG</u>	<u>A203409</u>	<u>001</u>	Aug 31, 2015
<u>AB</u>			<u>2MG</u>	<u>A203409</u>	<u>002</u>	Aug 31, 2015
<u>AB</u>		UNICHEM	<u>1MG</u>	<u>A205399</u>	<u>001</u>	Aug 05, 2020
<u>AB</u>			<u>2MG</u>	<u>A205399</u>	<u>002</u>	Aug 05, 2020
<u>AB</u>		UNIQUE	<u>1MG</u>	<u>A204721</u>	<u>001</u>	Jan 24, 2020
<u>AB</u>			<u>2MG</u>	<u>A204721</u>	<u>002</u>	Jan 24, 2020

TOLVAPTAN

TABLET;ORAL

SAMSCA

<u>AB</u>	+	OTSUKA	<u>15MG</u>	<u>N022275</u>	<u>001</u>	May 19, 2009
<u>AB</u>	+	!	<u>30MG</u>	<u>N022275</u>	<u>002</u>	May 19, 2009

TOLVAPTAN

<u>AB</u>		ALKEM LABS LTD	<u>15MG</u>	<u>A211891</u>	<u>003</u>	Sep 06, 2022
<u>AB</u>			<u>30MG</u>	<u>A211891</u>	<u>001</u>	May 19, 2020
<u>AB</u>		APOTEX	<u>15MG</u>	<u>A207605</u>	<u>002</u>	Sep 06, 2022
<u>AB</u>			<u>30MG</u>	<u>A207605</u>	<u>001</u>	May 19, 2020
<u>AB</u>		ENDO OPERATIONS	<u>15MG</u>	<u>A206119</u>	<u>001</u>	Feb 15, 2022
<u>AB</u>			<u>30MG</u>	<u>A206119</u>	<u>002</u>	Feb 15, 2022
<u>AB</u>		HETERO LABS LTD V	<u>15MG</u>	<u>A205646</u>	<u>002</u>	Sep 06, 2022
<u>AB</u>			<u>30MG</u>	<u>A205646</u>	<u>001</u>	Jul 16, 2021
<u>AB</u>		MSN	<u>15MG</u>	<u>A216949</u>	<u>001</u>	Jun 21, 2023
<u>AB</u>			<u>30MG</u>	<u>A216949</u>	<u>002</u>	Jun 21, 2023

JYNARQUE

+	OTSUKA	15MG	N204441	001	Apr 23, 2018
---	--------	------	---------	-----	--------------

+		30MG	N204441	002	Apr 23, 2018
---	--	------	---------	-----	--------------

+	!	45MG	N204441	003	Apr 23, 2018
---	---	------	---------	-----	--------------

+		60MG	N204441	004	Apr 23, 2018
---	--	------	---------	-----	--------------

+		90MG	N204441	005	Apr 23, 2018
---	--	------	---------	-----	--------------

TOLVAPTAN

	ALKEM LABS LTD	60MG	A211891	002	May 19, 2020
--	----------------	------	---------	-----	--------------

TOPIRAMATE

CAPSULE;ORAL

TOPAMAX

<u>AB</u>	+	JANSSEN PHARMS	<u>15MG</u>	<u>N020844</u>	<u>001</u>	Oct 26, 1998
<u>AB</u>	+	!	<u>25MG</u>	<u>N020844</u>	<u>002</u>	Oct 26, 1998

TOPIRAMATE

<u>AB</u>		ALKEM LABS LTD	<u>15MG</u>	<u>A218482</u>	<u>001</u>	Jun 13, 2024
<u>AB</u>			<u>25MG</u>	<u>A218482</u>	<u>002</u>	Jun 13, 2024
<u>AB</u>		AUROBINDO PHARMA LTD	<u>15MG</u>	<u>A215449</u>	<u>001</u>	Sep 27, 2023
<u>AB</u>			<u>25MG</u>	<u>A215449</u>	<u>002</u>	Sep 27, 2023
<u>AB</u>		GLENMARK PHARMS LTD	<u>15MG</u>	<u>A217869</u>	<u>001</u>	Jul 16, 2024
<u>AB</u>			<u>25MG</u>	<u>A217869</u>	<u>002</u>	Jul 16, 2024
<u>AB</u>		SENORES PHARMS	<u>15MG</u>	<u>A218642</u>	<u>001</u>	Oct 10, 2024
<u>AB</u>			<u>25MG</u>	<u>A218642</u>	<u>002</u>	Oct 10, 2024
<u>AB</u>		STRIDES PHARMA	<u>15MG</u>	<u>A078418</u>	<u>001</u>	Oct 14, 2009
<u>AB</u>			<u>25MG</u>	<u>A078418</u>	<u>002</u>	Oct 14, 2009
<u>AB</u>		TEVA	<u>15MG</u>	<u>A076575</u>	<u>001</u>	Apr 17, 2009
<u>AB</u>			<u>25MG</u>	<u>A076575</u>	<u>002</u>	Apr 17, 2009
<u>AB</u>		TWI PHARMS	<u>15MG</u>	<u>A217694</u>	<u>001</u>	Dec 05, 2023
<u>AB</u>			<u>25MG</u>	<u>A217694</u>	<u>002</u>	Dec 05, 2023
<u>AB</u>		ZYDUS PHARMS USA INC	<u>15MG</u>	<u>A078877</u>	<u>001</u>	Oct 14, 2009
<u>AB</u>			<u>25MG</u>	<u>A078877</u>	<u>002</u>	Oct 14, 2009
			50MG	A078877	003	Oct 23, 2024

CAPSULE, EXTENDED RELEASE;ORAL

TOPIRAMATE

<u>AB1</u>		ACTAVIS LABS FL	<u>25MG</u>	<u>A206210</u>	<u>001</u>	Mar 10, 2023
<u>AB1</u>			<u>50MG</u>	<u>A206210</u>	<u>002</u>	Mar 10, 2023

PRESCRIPTION DRUG PRODUCT LISTTOPIRAMATE

CAPSULE, EXTENDED RELEASE;ORAL

TOPIRAMATE

<u>AB1</u>		<u>100MG</u>	<u>A206210 003</u>	Mar 10, 2023
<u>AB1</u>		<u>200MG</u>	<u>A206210 004</u>	Mar 10, 2023
<u>AB1</u>	ALKEM LABS LTD	<u>25MG</u>	<u>A217248 001</u>	Jun 20, 2023
<u>AB1</u>		<u>50MG</u>	<u>A217248 002</u>	Jun 20, 2023
<u>AB1</u>		<u>100MG</u>	<u>A217248 003</u>	Jun 20, 2023
<u>AB1</u>		<u>200MG</u>	<u>A217248 004</u>	Jun 20, 2023
<u>AB1</u>	DR REDDYS	<u>25MG</u>	<u>A217231 001</u>	Nov 17, 2023
<u>AB1</u>		<u>50MG</u>	<u>A217231 002</u>	Nov 17, 2023
<u>AB1</u>		<u>100MG</u>	<u>A217231 003</u>	Nov 17, 2023
<u>AB1</u>		<u>200MG</u>	<u>A217231 004</u>	Nov 17, 2023
<u>AB1</u>	ENDO OPERATIONS	<u>25MG</u>	<u>A205976 002</u>	May 04, 2023
<u>AB1</u>		<u>50MG</u>	<u>A205976 003</u>	May 04, 2023
<u>AB1</u>		<u>100MG</u>	<u>A205976 004</u>	May 04, 2023
<u>AB1</u>		<u>200MG</u>	<u>A205976 001</u>	Mar 01, 2023
<u>AB1</u>	LUPIN LTD	<u>25MG</u>	<u>A215561 001</u>	Jul 11, 2024
<u>AB1</u>		<u>50MG</u>	<u>A215561 002</u>	Jul 11, 2024
<u>AB1</u>		<u>100MG</u>	<u>A215561 003</u>	Jul 11, 2024
<u>AB1</u>		<u>200MG</u>	<u>A215561 004</u>	Jul 11, 2024
<u>AB1</u>	ZYDUS	<u>25MG</u>	<u>A207382 001</u>	Nov 24, 2017
<u>AB1</u>		<u>50MG</u>	<u>A207382 002</u>	Nov 24, 2017
<u>AB1</u>		<u>100MG</u>	<u>A207382 003</u>	Nov 24, 2017
<u>AB1</u>		<u>200MG</u>	<u>A207382 004</u>	Oct 30, 2023

TROKENDI XR

<u>AB1</u>	+	SUPERNUS PHARMS	<u>25MG</u>	<u>N201635 001</u>	Aug 16, 2013
<u>AB1</u>	+		<u>50MG</u>	<u>N201635 002</u>	Aug 16, 2013
<u>AB1</u>	+		<u>100MG</u>	<u>N201635 003</u>	Aug 16, 2013
<u>AB1</u>	+	!	<u>200MG</u>	<u>N201635 004</u>	Aug 16, 2013

OUDEXY XR

<u>AB2</u>	+	UPSHER SMITH LABS	<u>25MG</u>	<u>N205122 001</u>	Mar 11, 2014
<u>AB2</u>	+		<u>50MG</u>	<u>N205122 002</u>	Mar 11, 2014
<u>AB2</u>	+		<u>100MG</u>	<u>N205122 003</u>	Mar 11, 2014
<u>AB2</u>	+		<u>150MG</u>	<u>N205122 004</u>	Mar 11, 2014
<u>AB2</u>	+	!	<u>200MG</u>	<u>N205122 005</u>	Mar 11, 2014

TOPIRAMATE

<u>AB2</u>		AMTA	<u>25MG</u>	<u>A218695 001</u>	Dec 03, 2024
<u>AB2</u>			<u>50MG</u>	<u>A218695 002</u>	Dec 03, 2024
<u>AB2</u>			<u>100MG</u>	<u>A218695 003</u>	Dec 03, 2024
<u>AB2</u>			<u>150MG</u>	<u>A218695 004</u>	Dec 03, 2024
<u>AB2</u>			<u>200MG</u>	<u>A218695 005</u>	Dec 03, 2024
<u>AB2</u>		GLENMARK PHARMS LTD	<u>25MG</u>	<u>A210278 001</u>	Feb 01, 2021
<u>AB2</u>			<u>50MG</u>	<u>A210278 002</u>	Feb 01, 2021
<u>AB2</u>			<u>100MG</u>	<u>A210278 003</u>	Feb 01, 2021
<u>AB2</u>			<u>150MG</u>	<u>A210278 004</u>	Feb 01, 2021
<u>AB2</u>			<u>200MG</u>	<u>A210278 005</u>	Feb 01, 2021
<u>AB2</u>		XIAMEN LP PHARM CO	<u>25MG</u>	<u>A215638 001</u>	Nov 05, 2024
<u>AB2</u>			<u>50MG</u>	<u>A215638 002</u>	Nov 05, 2024
<u>AB2</u>			<u>100MG</u>	<u>A215638 003</u>	Nov 05, 2024
<u>AB2</u>			<u>150MG</u>	<u>A215638 004</u>	Nov 05, 2024
<u>AB2</u>			<u>200MG</u>	<u>A215638 005</u>	Nov 05, 2024
<u>AB2</u>		ZYDUS	<u>25MG</u>	<u>A208949 001</u>	Nov 29, 2022
<u>AB2</u>			<u>50MG</u>	<u>A208949 002</u>	Nov 29, 2022
<u>AB2</u>			<u>100MG</u>	<u>A208949 003</u>	Nov 29, 2022
<u>AB2</u>			<u>150MG</u>	<u>A208949 004</u>	Nov 29, 2022
<u>AB2</u>			<u>200MG</u>	<u>A208949 005</u>	Nov 29, 2022

SOLUTION;ORAL

EPRONTIA

<u>AB</u>	+	AZURITY	<u>25MG/ML</u>	<u>N214679 001</u>	Nov 05, 2021
-----------	---	---------	----------------	--------------------	--------------

TOPIRAMATE

<u>AB</u>		ALKEM LABS LTD	<u>25MG/ML</u>	<u>A217795 001</u>	Oct 31, 2024
-----------	--	----------------	----------------	--------------------	--------------

TABLET;ORAL

TOPAMAX

<u>AB</u>	+	JANSSEN PHARMS	<u>25MG</u>	<u>N020505 004</u>	Dec 24, 1996
<u>AB</u>	+		<u>50MG</u>	<u>N020505 005</u>	Dec 24, 1996
<u>AB</u>	+	!	<u>100MG</u>	<u>N020505 001</u>	Dec 24, 1996
<u>AB</u>	+		<u>200MG</u>	<u>N020505 002</u>	Dec 24, 1996

TOPIRAMATE

<u>AB</u>		ACCORD HLTHCARE	<u>25MG</u>	<u>A076311 001</u>	Mar 27, 2009
<u>AB</u>			<u>50MG</u>	<u>A076311 002</u>	Mar 27, 2009
<u>AB</u>			<u>100MG</u>	<u>A076311 003</u>	Mar 27, 2009

PRESCRIPTION DRUG PRODUCT LIST

TOPIRAMATE

TABLET; ORAL

TOPIRAMATE

<u>AB</u>		<u>200MG</u>	<u>A076311</u>	<u>004</u>	Mar 27, 2009
<u>AB</u>	ASCENT PHARMS INC	<u>25MG</u>	<u>A215414</u>	<u>001</u>	Aug 26, 2021
<u>AB</u>		<u>50MG</u>	<u>A215414</u>	<u>002</u>	Aug 26, 2021
<u>AB</u>		<u>100MG</u>	<u>A215414</u>	<u>003</u>	Aug 26, 2021
<u>AB</u>		<u>200MG</u>	<u>A215414</u>	<u>004</u>	Aug 26, 2021
<u>AB</u>	AUROBINDO PHARMA	<u>25MG</u>	<u>A078462</u>	<u>001</u>	Mar 27, 2009
<u>AB</u>		<u>50MG</u>	<u>A078462</u>	<u>002</u>	Mar 27, 2009
<u>AB</u>		<u>100MG</u>	<u>A078462</u>	<u>003</u>	Mar 27, 2009
<u>AB</u>		<u>200MG</u>	<u>A078462</u>	<u>004</u>	Mar 27, 2009
<u>AB</u>	CIPLA LTD	<u>25MG</u>	<u>A076343</u>	<u>001</u>	Mar 27, 2009
<u>AB</u>		<u>50MG</u>	<u>A076343</u>	<u>002</u>	Mar 27, 2009
<u>AB</u>		<u>100MG</u>	<u>A076343</u>	<u>003</u>	Mar 27, 2009
<u>AB</u>		<u>200MG</u>	<u>A076343</u>	<u>004</u>	Mar 27, 2009
<u>AB</u>	GLENMARK PHARMS LTD	<u>25MG</u>	<u>A077627</u>	<u>001</u>	Mar 27, 2009
<u>AB</u>		<u>50MG</u>	<u>A077627</u>	<u>002</u>	Mar 27, 2009
<u>AB</u>		<u>100MG</u>	<u>A077627</u>	<u>003</u>	Mar 27, 2009
<u>AB</u>		<u>200MG</u>	<u>A077627</u>	<u>004</u>	Mar 27, 2009
<u>AB</u>	INVAGEN PHARMS	<u>25MG</u>	<u>A079162</u>	<u>001</u>	Mar 27, 2009
<u>AB</u>		<u>50MG</u>	<u>A079162</u>	<u>002</u>	Mar 27, 2009
<u>AB</u>		<u>100MG</u>	<u>A079162</u>	<u>003</u>	Mar 27, 2009
<u>AB</u>		<u>200MG</u>	<u>A079162</u>	<u>004</u>	Mar 27, 2009
<u>AB</u>	SUN PHARM	<u>25MG</u>	<u>A090278</u>	<u>001</u>	Mar 27, 2009
<u>AB</u>		<u>50MG</u>	<u>A090278</u>	<u>002</u>	Mar 27, 2009
<u>AB</u>		<u>100MG</u>	<u>A090278</u>	<u>003</u>	Mar 27, 2009
<u>AB</u>		<u>200MG</u>	<u>A090278</u>	<u>004</u>	Mar 27, 2009
<u>AB</u>	UNICHEM LABS LTD	<u>25MG</u>	<u>A090162</u>	<u>001</u>	Mar 27, 2009
<u>AB</u>		<u>50MG</u>	<u>A090162</u>	<u>002</u>	Mar 27, 2009
<u>AB</u>		<u>100MG</u>	<u>A090162</u>	<u>003</u>	Mar 27, 2009
<u>AB</u>		<u>200MG</u>	<u>A090162</u>	<u>004</u>	Feb 19, 2013
<u>AB</u>	VIWIT PHARM	<u>25MG</u>	<u>A077733</u>	<u>001</u>	Mar 27, 2009
<u>AB</u>		<u>50MG</u>	<u>A077733</u>	<u>002</u>	Mar 27, 2009
<u>AB</u>		<u>100MG</u>	<u>A077733</u>	<u>003</u>	Mar 27, 2009
<u>AB</u>		<u>200MG</u>	<u>A077733</u>	<u>004</u>	Mar 27, 2009
<u>AB</u>	ZYDUS PHARMS USA INC	<u>25MG</u>	<u>A078235</u>	<u>001</u>	Mar 27, 2009
<u>AB</u>		<u>50MG</u>	<u>A078235</u>	<u>002</u>	Mar 27, 2009
<u>AB</u>		<u>100MG</u>	<u>A078235</u>	<u>003</u>	Mar 27, 2009
<u>AB</u>		<u>200MG</u>	<u>A078235</u>	<u>004</u>	Mar 27, 2009

TOPOTECAN HYDROCHLORIDE

CAPSULE; ORAL

HYCAMTIN

+ SANDOZ

EQ 0.25MG BASE

N020981 001 Oct 11, 2007

+!

EQ 1MG BASE

N020981 002 Oct 11, 2007

INJECTABLE; INJECTION

HYCAMTIN

<u>AP</u>	<u>+!</u> SANDOZ	<u>EQ 4MG BASE/VIAL</u>	<u>N020671</u>	<u>001</u>	May 28, 1996
	<u>TOPOTECAN HYDROCHLORIDE</u>				
<u>AP</u>	ACCORD HLTHCARE	<u>EQ 4MG BASE/VIAL</u>	<u>A202351</u>	<u>001</u>	Jun 26, 2013
<u>AP</u>	ACTAVIS TOTOWA	<u>EQ 4MG BASE/VIAL</u>	<u>A090620</u>	<u>001</u>	Dec 02, 2010
<u>AP</u>	DR REDDYS LABS LTD	<u>EQ 4MG BASE/VIAL</u>	<u>A201191</u>	<u>001</u>	Mar 09, 2011
<u>AP</u>	FRESENIUS KABI USA	<u>EQ 4MG BASE/VIAL</u>	<u>A091089</u>	<u>001</u>	Nov 29, 2010
<u>AP</u>	NOVAST LABS	<u>EQ 4MG BASE/VIAL</u>	<u>A206962</u>	<u>001</u>	Nov 30, 2016
<u>AP</u>	SAGENT PHARMS	<u>EQ 4MG BASE/VIAL</u>	<u>A091284</u>	<u>001</u>	Jan 26, 2011
<u>AP</u>	TEYRO LABS	<u>EQ 4MG BASE/VIAL</u>	<u>A091199</u>	<u>001</u>	Dec 01, 2010

SOLUTION; INTRAVENOUS

TOPOTECAN HYDROCHLORIDE

<u>AP</u>	ACCORD HLTHCARE	<u>EQ 4MG BASE/4ML (EQ 1MG BASE/ML)</u>	<u>A204406</u>	<u>002</u>	Jul 06, 2017
<u>AP</u>	<u>+!</u> HOSPIRA INC	<u>EQ 4MG BASE/4ML (EQ 1MG BASE/ML)</u>	<u>N200582</u>	<u>001</u>	Feb 02, 2011
	ACCORD HLTHCARE	EQ 1MG BASE/ML (EQ 1MG BASE/ML)	A204406	001	Jul 06, 2017

TOREMIFENE CITRATE

TABLET; ORAL

TOREMIFENE CITRATE

<u>AB</u>	MSN	<u>EQ 60MG BASE</u>	<u>A212818</u>	<u>001</u>	Aug 18, 2020
<u>AB</u>	<u>!</u> RISING	<u>EQ 60MG BASE</u>	<u>A208813</u>	<u>001</u>	Dec 04, 2018

PRESCRIPTION DRUG PRODUCT LIST

TORSEMIDE

TABLET; ORAL

TORSEMIDE

<u>AB</u>	AUROBINDO PHARMA	<u>5MG</u>	<u>A078249</u>	<u>001</u>	Oct 17, 2007
<u>AB</u>		<u>10MG</u>	<u>A078249</u>	<u>002</u>	Oct 17, 2007
<u>AB</u>		<u>20MG</u>	<u>A078249</u>	<u>003</u>	Oct 17, 2007
<u>AB</u>		<u>100MG</u>	<u>A078249</u>	<u>004</u>	Oct 17, 2007
<u>AB</u>	HERITAGE	<u>5MG</u>	<u>A076894</u>	<u>001</u>	May 31, 2005
<u>AB</u>		<u>10MG</u>	<u>A076894</u>	<u>002</u>	May 31, 2005
<u>AB</u>		<u>20MG</u>	<u>A076894</u>	<u>003</u>	May 31, 2005
<u>AB</u>		<u>100MG</u>	<u>A076894</u>	<u>004</u>	May 31, 2005
<u>AB</u>	HETERO LABS LTD III	<u>5MG</u>	<u>A079234</u>	<u>001</u>	Jan 27, 2009
<u>AB</u>		<u>10MG</u>	<u>A079234</u>	<u>002</u>	Jan 27, 2009
<u>AB</u>		<u>20MG</u>	<u>A079234</u>	<u>003</u>	Jan 27, 2009
<u>AB</u>		<u>100MG</u>	<u>A079234</u>	<u>004</u>	Jan 27, 2009
<u>AB</u>	HIKMA	<u>5MG</u>	<u>A076943</u>	<u>001</u>	Mar 01, 2005
<u>AB</u>		<u>10MG</u>	<u>A076943</u>	<u>002</u>	Mar 01, 2005
<u>AB</u>		<u>20MG</u>	<u>A076943</u>	<u>003</u>	Mar 01, 2005
<u>AB</u>	PLIVA PHARM IND	<u>5MG</u>	<u>A076346</u>	<u>001</u>	May 30, 2003
<u>AB</u>		<u>10MG</u>	<u>A076346</u>	<u>002</u>	May 30, 2003
<u>AB</u>	!	<u>20MG</u>	<u>A076346</u>	<u>003</u>	May 30, 2003
<u>AB</u>		<u>100MG</u>	<u>A076346</u>	<u>004</u>	Oct 19, 2004
<u>AB</u>	STRIDES PHARMA	<u>5MG</u>	<u>A076226</u>	<u>001</u>	May 27, 2003
<u>AB</u>		<u>5MG</u>	<u>A090613</u>	<u>001</u>	Mar 22, 2011
<u>AB</u>		<u>10MG</u>	<u>A076226</u>	<u>002</u>	May 27, 2003
<u>AB</u>		<u>10MG</u>	<u>A090613</u>	<u>002</u>	Mar 22, 2011
<u>AB</u>		<u>20MG</u>	<u>A076226</u>	<u>003</u>	May 27, 2003
<u>AB</u>		<u>20MG</u>	<u>A090613</u>	<u>003</u>	Mar 22, 2011
<u>AB</u>		<u>100MG</u>	<u>A076226</u>	<u>004</u>	May 27, 2003
<u>AB</u>		<u>100MG</u>	<u>A090613</u>	<u>004</u>	Mar 22, 2011
	SOAANZ				
	+ SARFE PHARMS	40MG	N213218	003	Nov 17, 2021

TOVORAFENIB

FOR SUSPENSION; ORAL

OJEMDA

+! DAY ONE BIOPHARMS 25MG/ML N218033 001 Apr 23, 2024

TABLET; ORAL

OJEMDA

+! DAY ONE BIOPHARMS 100MG N217700 001 Apr 23, 2024

TRABECTEDIN

POWDER; INTRAVENOUS

YONDELIS

+! JANSSEN PRODS 1MG/VIAL N207953 001 Oct 23, 2015

TRAMADOL HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

CONZIP

+! CIPHER PHARMS INC 100MG N022370 001 May 07, 2010

+ 200MG N022370 002 May 07, 2010

+ 300MG N022370 003 May 07, 2010

TABLET; ORAL

TRAMADOL HYDROCHLORIDE

<u>AB</u>	AMNEAL PHARMS	<u>50MG</u>	<u>A076003</u>	<u>001</u>	Jun 20, 2002
<u>AB</u>	APOTEX	<u>50MG</u>	<u>A075981</u>	<u>001</u>	Jul 10, 2002
<u>AB</u>	AUROBINDO PHARMA LTD	<u>50MG</u>	<u>A203494</u>	<u>001</u>	Mar 31, 2014
<u>AB</u>	CSPC OUYI PHARM CO	<u>50MG</u>	<u>A091498</u>	<u>001</u>	Mar 29, 2013
<u>AB</u>	GRAVITI PHARMS	<u>100MG</u>	<u>A075968</u>	<u>002</u>	Aug 02, 2024
<u>AB</u>	MERRO PHARM USA	<u>50MG</u>	<u>A206706</u>	<u>001</u>	Jul 02, 2019
<u>AB</u>	RUBICON	<u>50MG</u>	<u>A208708</u>	<u>001</u>	Jun 28, 2019
<u>AB</u>	!	<u>100MG</u>	<u>A208708</u>	<u>002</u>	Jun 28, 2019
<u>AB</u>	STRIDES PHARMA	<u>50MG</u>	<u>A202075</u>	<u>001</u>	Nov 28, 2011
<u>AB</u>	SUN PHARM INDS INC	<u>50MG</u>	<u>A075964</u>	<u>001</u>	Jun 19, 2002
<u>AB</u>	TEVA	<u>50MG</u>	<u>A075977</u>	<u>001</u>	Jun 19, 2002
<u>AB</u>	UNICHEM	<u>50MG</u>	<u>A211825</u>	<u>001</u>	Aug 09, 2019
<u>AB</u>	ZYDUS PHARMS USA INC	<u>50MG</u>	<u>A090404</u>	<u>001</u>	Jan 31, 2011
	RUBICON	25MG	A208708	003	Nov 27, 2023
		75MG	A208708	004	Apr 04, 2024
	TABLET, EXTENDED RELEASE; ORAL				
	TRAMADOL HYDROCHLORIDE				
	! LUPIN LTD	100MG	A200503	001	Aug 29, 2011

PRESCRIPTION DRUG PRODUCT LISTTRAMADOL HYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL
TRAMADOL HYDROCHLORIDE

200MG A200503 002 Aug 29, 2011
300MG A200503 003 Aug 29, 2011

TRAMETINIB DIMETHYL SULFOXIDE

SOLUTION;ORAL
MEKINIST

+! NOVARTIS EQ 0.05MG BASE/ML N217513 001 Mar 16, 2023

TABLET;ORAL

MEKINIST

+ NOVARTIS EQ 0.5MG N204114 001 May 29, 2013
+! EQ 2MG N204114 003 May 29, 2013

TRANDOLAPRIL

TABLET;ORAL

TRANDOLAPRIL

<u>AB</u>	AUROBINDO PHARMA	<u>1MG</u>	<u>A078438 001</u>	Jun 12, 2007
<u>AB</u>		<u>2MG</u>	<u>A078438 002</u>	Jun 12, 2007
<u>AB</u>		<u>4MG</u>	<u>A078438 003</u>	Jun 12, 2007
<u>AB</u>	EPIC PHARMA	<u>1MG</u>	<u>A078508 003</u>	Jun 18, 2008
<u>AB</u>		<u>2MG</u>	<u>A078508 001</u>	Jun 18, 2008
<u>AB</u>		<u>4MG</u>	<u>A078508 002</u>	Jun 18, 2008
<u>AB</u>	LUPIN	<u>1MG</u>	<u>A077522 001</u>	Jun 12, 2007
<u>AB</u>		<u>2MG</u>	<u>A077522 002</u>	Jun 12, 2007
<u>AB</u>	!	<u>4MG</u>	<u>A077522 003</u>	Jun 12, 2007

TRANDOLAPRIL; VERAPAMIL HYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL

TRANDOLAPRIL AND VERAPAMIL HYDROCHLORIDE

GLENMARK PHARMS LTD 1MG;240MG A079135 004 Aug 30, 2010
2MG;180MG A079135 001 May 26, 2010
2MG;240MG A079135 002 May 26, 2010
! 4MG;240MG A079135 003 May 05, 2010

TRANEXAMIC ACID

INJECTABLE; INJECTION

CYKLOKAPRON

AP +! PFIZER 100MG/ML N019281 001 Dec 30, 1986

TRANEXAMIC ACID

<u>AP</u>	AM REGENT	<u>100MG/ML</u>	<u>A201885 001</u>	Aug 10, 2011
<u>AP</u>	AMNEAL PHARMS CO	<u>100MG/ML</u>	<u>A208840 001</u>	Feb 28, 2017
<u>AP</u>	APOTEX	<u>100MG/ML</u>	<u>A209860 001</u>	Jan 14, 2020
<u>AP</u>	AVET LIFESCIENCES	<u>100MG/ML</u>	<u>A203521 001</u>	Aug 12, 2014
<u>AP</u>	CAPLIN	<u>100MG/ML</u>	<u>A212360 001</u>	Jul 17, 2019
<u>AP</u>	EPIC PHARMA LLC	<u>100MG/ML</u>	<u>A202373 001</u>	Nov 17, 2011
<u>AP</u>	EUGIA PHARMA	<u>100MG/ML</u>	<u>A205035 001</u>	Jan 14, 2016
<u>AP</u>	FRESENIUS KABI USA	<u>100MG/ML</u>	<u>A091596 001</u>	Mar 02, 2012
<u>AP</u>	GLAND	<u>100MG/ML</u>	<u>A207239 001</u>	Feb 13, 2017
<u>AP</u>	HERITAGE	<u>100MG/ML</u>	<u>A202436 001</u>	Feb 11, 2014
<u>AP</u>	MICRO LABS	<u>100MG/ML</u>	<u>A206713 001</u>	Jun 27, 2017
<u>AP</u>	MYLAN INSTITUTIONAL	<u>100MG/ML</u>	<u>A091657 001</u>	Nov 03, 2011
<u>AP</u>	PROVEPHARM SAS	<u>100MG/ML</u>	<u>A212676 001</u>	Jul 17, 2019
<u>AP</u>	XGEN PHARMS	<u>100MG/ML</u>	<u>A201580 001</u>	Jun 14, 2013

SOLUTION; INTRAVENOUS

TRANEXAMIC ACID

<u>AP</u>	AMNEAL	<u>1GM/100ML (10MG/ML)</u>	<u>A217155 001</u>	Oct 16, 2023
<u>AP</u>	+! EXELA PHARMA	<u>1GM/100ML (10MG/ML)</u>	<u>N212020 001</u>	Apr 15, 2019
<u>AP</u>	FRESENIUS KABI USA	<u>1GM/100ML (10MG/ML)</u>	<u>A218242 001</u>	Aug 15, 2024
<u>AP</u>	GLAND PHARMA LTD	<u>1GM/100ML (10MG/ML)</u>	<u>A218599 001</u>	Apr 16, 2024
<u>AP</u>	NEXUS	<u>1GM/100ML (10MG/ML)</u>	<u>A216877 001</u>	Apr 16, 2024

TABLET;ORAL

LYSTEDA

AB +! AMRING PHARMS 650MG N022430 001 Nov 13, 2009

TRANEXAMIC ACID

<u>AB</u>	ACTAVIS LABS FL INC	<u>650MG</u>	<u>A202093 001</u>	Dec 27, 2012
<u>AB</u>	ANI PHARMS	<u>650MG</u>	<u>A203256 001</u>	Jul 25, 2016
<u>AB</u>	AUROBINDO PHARMA USA	<u>650MG</u>	<u>A205133 001</u>	Sep 21, 2015
<u>AB</u>	RUBICON	<u>650MG</u>	<u>A218320 001</u>	Jun 11, 2024

PRESCRIPTION DRUG PRODUCT LIST

TRANLYCYPROMINE SULFATE

TABLET; ORAL

PARNATE

AB +! ADVANZ PHARMA **EQ 10MG BASE** **N012342 003** Aug 16, 1985

TRANLYCYPROMINE SULFATE

AB CROSSMEDIKA SA **EQ 10MG BASE** **A213503 001** Jun 27, 2022
AB NOVITIUM PHARMA **EQ 10MG BASE** **A206856 001** Apr 17, 2018
AB STRIDES PHARMA **EQ 10MG BASE** **A040640 001** Jun 29, 2006

TRAVOPROST

IMPLANT; INTRACAMERAL

IDOSE TR

+! GLAUKOS 75MCG N218010 001 Dec 13, 2023

SOLUTION/DROPS; OPHTHALMIC

TRAVATAN Z

AT +! SANDOZ **0.004%** **N021994 001** Sep 21, 2006

TRAVOPROST

AT APOTEX **0.004%** **A203431 001** Jul 10, 2015
AT GLAND PHARMA LTD **0.004%** **A218159 001** Jul 12, 2024
AT MICRO LABS **0.004%** **A203767 001** Mar 19, 2021
AT MYLAN **0.004%** **A205050 001** Jul 07, 2017
AT SOMERSET THERAPS LLC **0.004%** **A217228 001** Sep 23, 2024
AT1 ALEMBIC **0.004%** **A210458 001** Dec 20, 2019
AT1 ! CHARTWELL RX **0.004%** **A091340 001** Mar 01, 2013

TRAZODONE HYDROCHLORIDE

SOLUTION; ORAL

RALDESY

+! KAMAT 10MG/ML N218637 001 Nov 26, 2024

TABLET; ORAL

TRAZODONE HYDROCHLORIDE

AB ACCORD HLTHCARE **50MG** **A206923 001** Sep 08, 2017
AB **100MG** **A206923 002** Sep 08, 2017
AB **150MG** **A206923 003** Sep 08, 2017
AB **300MG** **A206923 004** Sep 08, 2017
AB APOTEX **50MG** **A071258 001** Mar 25, 1987
AB ! APOTEX INC **100MG** **A071196 001** Mar 25, 1987
AB **150MG** **A071196 002** Apr 26, 1999
AB **300MG** **A071196 003** Apr 26, 1999
AB AUROBINDO PHARMA **50MG** **A204852 001** Feb 05, 2020
AB **100MG** **A204852 002** Feb 05, 2020
AB **150MG** **A204852 003** Feb 05, 2020
AB **300MG** **A204852 004** Feb 05, 2020
AB CROSSMEDIKA SA **50MG** **A211116 001** May 20, 2024
AB **100MG** **A211116 002** May 20, 2024
AB **150MG** **A211116 003** May 20, 2024
AB **300MG** **A211116 004** May 20, 2024
AB GRANULES **50MG** **A218988 001** Aug 08, 2024
AB **100MG** **A218988 002** Aug 08, 2024
AB **150MG** **A218988 003** Aug 08, 2024
AB **300MG** **A218988 004** Aug 08, 2024
AB GRAVITI PHARMS **50MG** **A217740 001** Aug 27, 2024
AB **100MG** **A217740 002** Aug 27, 2024
AB **150MG** **A217740 003** Aug 27, 2024
AB **300MG** **A217740 004** Aug 27, 2024
AB OXFORD PHARMS **50MG** **A072192 001** Feb 02, 1989
AB **100MG** **A072193 001** Feb 02, 1989
AB PLIVA **150MG** **A071525 001** Mar 09, 1988
AB SUN PHARM INDUSTRIES **50MG** **A073137 002** Mar 24, 1993
AB **100MG** **A073137 001** Mar 24, 1993
AB **150MG** **A073137 003** Dec 22, 1995
AB TEVA PHARMS USA **50MG** **A071523 001** Dec 11, 1987
AB **100MG** **A071524 001** Dec 11, 1987
AB TORRENT **50MG** **A202180 001** Nov 27, 2013
AB **100MG** **A202180 002** Nov 27, 2013
AB **150MG** **A202180 003** Nov 27, 2013
AB **300MG** **A202180 004** Nov 27, 2013
AB ZYDUS PHARMS **50MG** **A205253 001** Oct 10, 2017
AB **100MG** **A205253 002** Oct 10, 2017
AB **150MG** **A205253 003** Oct 10, 2017
AB **300MG** **A205253 004** Oct 10, 2017

PRESCRIPTION DRUG PRODUCT LIST

TREPROSTINIL

INJECTABLE; INTRAVENOUS, SUBCUTANEOUS

REMODULIN

<u>AP</u>	<u>+</u> !	UNITED THERAP	<u>1MG/ML</u>	<u>N021272</u>	<u>001</u>	May 21, 2002
<u>AP</u>	<u>+</u> !		<u>2.5MG/ML</u>	<u>N021272</u>	<u>002</u>	May 21, 2002
<u>AP</u>	<u>+</u> !		<u>5MG/ML</u>	<u>N021272</u>	<u>003</u>	May 21, 2002
<u>AP</u>	<u>+</u> !		<u>10MG/ML</u>	<u>N021272</u>	<u>004</u>	May 21, 2002

TREPROSTINIL

<u>AP</u>		ALEMBIC GLOBAL	<u>1MG/ML</u>	<u>A211574</u>	<u>001</u>	Feb 11, 2021
<u>AP</u>			<u>2.5MG/ML</u>	<u>A211574</u>	<u>002</u>	Feb 11, 2021
<u>AP</u>			<u>5MG/ML</u>	<u>A211574</u>	<u>003</u>	Feb 11, 2021
<u>AP</u>			<u>10MG/ML</u>	<u>A211574</u>	<u>004</u>	Feb 11, 2021
<u>AP</u>		DR REDDYS	<u>1MG/ML</u>	<u>A210214</u>	<u>001</u>	May 22, 2020
<u>AP</u>			<u>2.5MG/ML</u>	<u>A210214</u>	<u>002</u>	May 22, 2020
<u>AP</u>			<u>5MG/ML</u>	<u>A210214</u>	<u>003</u>	May 22, 2020
<u>AP</u>			<u>10MG/ML</u>	<u>A210214</u>	<u>004</u>	May 22, 2020
<u>AP</u>		ENDO OPERATIONS	<u>1MG/ML</u>	<u>A209382</u>	<u>001</u>	Sep 24, 2019
<u>AP</u>			<u>2.5MG/ML</u>	<u>A209382</u>	<u>002</u>	Sep 24, 2019
<u>AP</u>			<u>5MG/ML</u>	<u>A209382</u>	<u>003</u>	Sep 24, 2019
<u>AP</u>			<u>10MG/ML</u>	<u>A209382</u>	<u>004</u>	Sep 24, 2019
<u>AP</u>		SANDOZ	<u>1MG/ML</u>	<u>A203649</u>	<u>001</u>	Nov 30, 2017
<u>AP</u>			<u>2.5MG/ML</u>	<u>A203649</u>	<u>002</u>	Nov 30, 2017
<u>AP</u>			<u>5MG/ML</u>	<u>A203649</u>	<u>003</u>	Nov 30, 2017
<u>AP</u>			<u>10MG/ML</u>	<u>A203649</u>	<u>004</u>	Nov 30, 2017
<u>AP</u>		TEVA PHARMS USA	<u>1MG/ML</u>	<u>A206648</u>	<u>001</u>	Sep 26, 2019
<u>AP</u>			<u>2.5MG/ML</u>	<u>A206648</u>	<u>002</u>	Sep 26, 2019
<u>AP</u>			<u>5MG/ML</u>	<u>A206648</u>	<u>003</u>	Sep 26, 2019
<u>AP</u>			<u>10MG/ML</u>	<u>A206648</u>	<u>004</u>	Sep 26, 2019

REMODULIN

	<u>+</u> !	UNITED THERAP	0.1MG/ML	N021272	006	Sep 28, 2023
	<u>+</u> !		0.2MG/ML	N021272	007	Sep 28, 2023
	<u>+</u> !		0.4MG/ML	N021272	008	Sep 28, 2023
	<u>+</u> !		20MG/ML	N021272	005	Jul 30, 2021

POWDER; INHALATION

TYVASO DPI

	<u>+</u> !	UNITED THERAP	0.016MG/INH	N214324	001	May 23, 2022
	<u>+</u>		0.032MG/INH	N214324	002	May 23, 2022
	<u>+</u>		0.048MG/INH	N214324	003	May 23, 2022
	<u>+</u> !		0.064MG/INH	N214324	004	May 23, 2022
	<u>+</u>		0.08MG/INH	N214324	005	Oct 24, 2024

SOLUTION; INHALATION

TYVASO

	<u>+</u> !	UNITED THERAP	0.6MG/ML	N022387	001	Jul 30, 2009
--	------------	---------------	----------	---------	-----	--------------

TREPROSTINIL DIOLAMINE

TABLET, EXTENDED RELEASE; ORAL

ORENITRAM

	<u>+</u>	UNITED THERAP	EQ 0.125MG BASE	N203496	001	Dec 20, 2013
	<u>+</u>		EQ 0.25MG BASE	N203496	002	Dec 20, 2013
	<u>+</u> !		EQ 1MG BASE	N203496	003	Dec 20, 2013
	<u>+</u>		EQ 2.5MG BASE	N203496	004	Dec 20, 2013
	<u>+</u>		EQ 5MG BASE	N203496	005	Oct 07, 2016

TRETINOIN

CAPSULE; ORAL

TRETINOIN

<u>AB</u>	<u>!</u>	BARR LABS INC	<u>10MG</u>	<u>A077684</u>	<u>001</u>	Jun 22, 2007
<u>AB</u>		ENDO OPERATIONS	<u>10MG</u>	<u>A201687</u>	<u>001</u>	Oct 24, 2012
<u>AB</u>		GLENMARK PHARMS LTD	<u>10MG</u>	<u>A208279</u>	<u>001</u>	Dec 23, 2016

CREAM; TOPICAL

AVITA

<u>AB</u>		MYLAN PHARMS INC	<u>0.025%</u>	<u>N020404</u>	<u>003</u>	Jan 14, 1997
-----------	--	------------------	---------------	----------------	------------	--------------

RETIN-A

<u>AB</u>	<u>+</u> !	BAUSCH	<u>0.025%</u>	<u>N019049</u>	<u>001</u>	Sep 16, 1988
<u>AB</u>	<u>+</u> !	VALEANT PHARMS NORTH	<u>0.1%</u>	<u>N017340</u>	<u>001</u>	

TRETINOIN

<u>AB</u>		ALEMBIC	<u>0.1%</u>	<u>A217804</u>	<u>001</u>	Apr 29, 2024
<u>AB</u>		PADAGIS US	<u>0.025%</u>	<u>A075264</u>	<u>001</u>	Dec 24, 1998
<u>AB</u>			<u>0.1%</u>	<u>A075213</u>	<u>001</u>	Dec 24, 1998
<u>AB</u>		TARO	<u>0.025%</u>	<u>A215713</u>	<u>001</u>	Jan 05, 2023
<u>AB</u>			<u>0.1%</u>	<u>A211645</u>	<u>001</u>	Jan 22, 2019
<u>AB</u>		ZYDUS LIFESCIENCES	<u>0.1%</u>	<u>A218561</u>	<u>001</u>	Apr 25, 2024

PRESCRIPTION DRUG PRODUCT LIST

TRETINOIN

CREAM; TOPICAL

RETIN-A

<u>AB1</u>	+	VALEANT BERMUDA	<u>0.05%</u>	<u>N017522</u>	<u>001</u>	
------------	---	-----------------	--------------	----------------	------------	--

TRETINOIN

<u>AB1</u>		PADAGIS US	<u>0.05%</u>	<u>A075265</u>	<u>001</u>	Dec 24, 1998
------------	--	------------	--------------	----------------	------------	--------------

<u>AB1</u>		TARO	<u>0.05%</u>	<u>A211644</u>	<u>001</u>	Jan 25, 2019
------------	--	------	--------------	----------------	------------	--------------

RENOVA

+	VALEANT PHARMS	0.02%	N021108	001	Aug 31, 2000	
---	----------------	-------	---------	-----	--------------	--

NORTH

GEL; TOPICAL

ATRALIN

<u>AB</u>	+	DOW PHARM	<u>0.05%</u>	<u>N022070</u>	<u>001</u>	Jul 26, 2007
-----------	---	-----------	--------------	----------------	------------	--------------

RETIN-A

<u>AB</u>	+	VALEANT INTL	<u>0.01%</u>	<u>N017955</u>	<u>001</u>	
-----------	---	--------------	--------------	----------------	------------	--

<u>AB</u>	+		<u>0.025%</u>	<u>N017579</u>	<u>002</u>	
-----------	---	--	---------------	----------------	------------	--

RETIN-A-MICRO

<u>AB</u>	+	BAUSCH	<u>0.08%</u>	<u>N020475</u>	<u>003</u>	Jan 28, 2014
-----------	---	--------	--------------	----------------	------------	--------------

TRETINOIN

<u>AB</u>		MYLAN	<u>0.05%</u>	<u>A207955</u>	<u>001</u>	Aug 13, 2015
-----------	--	-------	--------------	----------------	------------	--------------

<u>AB</u>		PADAGIS US	<u>0.01%</u>	<u>A075589</u>	<u>001</u>	Jun 11, 2002
-----------	--	------------	--------------	----------------	------------	--------------

<u>AB</u>			<u>0.025%</u>	<u>A075529</u>	<u>001</u>	Feb 22, 2000
-----------	--	--	---------------	----------------	------------	--------------

TRETINOIN MICROSPHERE

<u>AB</u>		ENCUBE	<u>0.08%</u>	<u>A215609</u>	<u>001</u>	Aug 22, 2023
-----------	--	--------	--------------	----------------	------------	--------------

RETIN-A MICRO

+	BAUSCH	0.04%	N020475	002	May 10, 2002	
---	--------	-------	---------	-----	--------------	--

+		0.1%	N020475	001	Feb 07, 1997	
---	--	------	---------	-----	--------------	--

RETIN-A-MICRO

+	BAUSCH	0.06%	N020475	004	Oct 23, 2017	
---	--------	-------	---------	-----	--------------	--

LOTION; TOPICAL

ALTRENO

+	DOW PHARM	0.05%	N209353	001	Aug 23, 2018	
---	-----------	-------	---------	-----	--------------	--

TRIAMCINOLONE ACETONIDE

CREAM; TOPICAL

TRIAMCINOLONE ACETONIDE

<u>AT</u>		ALKEM LABS LTD	<u>0.025%</u>	<u>A207651</u>	<u>001</u>	Dec 26, 2017
-----------	--	----------------	---------------	----------------	------------	--------------

<u>AT</u>			<u>0.1%</u>	<u>A207651</u>	<u>002</u>	Dec 26, 2017
-----------	--	--	-------------	----------------	------------	--------------

<u>AT</u>			<u>0.5%</u>	<u>A207651</u>	<u>003</u>	Dec 26, 2017
-----------	--	--	-------------	----------------	------------	--------------

<u>AT</u>		CHARTWELL RX	<u>0.025%</u>	<u>A208763</u>	<u>001</u>	Feb 01, 2017
-----------	--	--------------	---------------	----------------	------------	--------------

<u>AT</u>			<u>0.1%</u>	<u>A208763</u>	<u>002</u>	Feb 01, 2017
-----------	--	--	-------------	----------------	------------	--------------

<u>AT</u>			<u>0.5%</u>	<u>A208763</u>	<u>003</u>	Feb 01, 2017
-----------	--	--	-------------	----------------	------------	--------------

<u>AT</u>		COSETTE	<u>0.025%</u>	<u>A089797</u>	<u>001</u>	May 31, 1991
-----------	--	---------	---------------	----------------	------------	--------------

<u>AT</u>			<u>0.1%</u>	<u>A089798</u>	<u>001</u>	May 31, 1991
-----------	--	--	-------------	----------------	------------	--------------

<u>AT</u>		ENCUBE	<u>0.1%</u>	<u>A208848</u>	<u>001</u>	Sep 18, 2017
-----------	--	--------	-------------	----------------	------------	--------------

<u>AT</u>	+	FOUGERA PHARMS	<u>0.025%</u>	<u>A085692</u>	<u>001</u>	
-----------	---	----------------	---------------	----------------	------------	--

<u>AT</u>	+		<u>0.1%</u>	<u>A085692</u>	<u>003</u>	
-----------	---	--	-------------	----------------	------------	--

<u>AT</u>	+		<u>0.5%</u>	<u>A085692</u>	<u>002</u>	
-----------	---	--	-------------	----------------	------------	--

<u>AT</u>		GLENMARK PHARMS LTD	<u>0.1%</u>	<u>A207117</u>	<u>001</u>	Aug 05, 2016
-----------	--	---------------------	-------------	----------------	------------	--------------

<u>AT</u>		MACLEODS PHARMS LTD	<u>0.025%</u>	<u>A209535</u>	<u>001</u>	May 18, 2018
-----------	--	---------------------	---------------	----------------	------------	--------------

<u>AT</u>			<u>0.1%</u>	<u>A209535</u>	<u>002</u>	May 18, 2018
-----------	--	--	-------------	----------------	------------	--------------

<u>AT</u>			<u>0.5%</u>	<u>A209535</u>	<u>003</u>	May 18, 2018
-----------	--	--	-------------	----------------	------------	--------------

<u>AT</u>		MICRO LABS	<u>0.025%</u>	<u>A040671</u>	<u>001</u>	Jun 09, 2006
-----------	--	------------	---------------	----------------	------------	--------------

<u>AT</u>			<u>0.1%</u>	<u>A040671</u>	<u>002</u>	Jun 09, 2006
-----------	--	--	-------------	----------------	------------	--------------

<u>AT</u>		PADAGIS US	<u>0.025%</u>	<u>A086413</u>	<u>002</u>	
-----------	--	------------	---------------	----------------	------------	--

<u>AT</u>			<u>0.1%</u>	<u>A086413</u>	<u>003</u>	
-----------	--	--	-------------	----------------	------------	--

<u>AT</u>			<u>0.5%</u>	<u>A086413</u>	<u>001</u>	
-----------	--	--	-------------	----------------	------------	--

<u>AT</u>		STRIDES PHARMA	<u>0.025%</u>	<u>A210346</u>	<u>001</u>	Feb 11, 2019
-----------	--	----------------	---------------	----------------	------------	--------------

<u>AT</u>			<u>0.1%</u>	<u>A210346</u>	<u>002</u>	Feb 11, 2019
-----------	--	--	-------------	----------------	------------	--------------

<u>AT</u>			<u>0.5%</u>	<u>A210346</u>	<u>003</u>	Feb 11, 2019
-----------	--	--	-------------	----------------	------------	--------------

<u>AT</u>		TARO	<u>0.1%</u>	<u>A040039</u>	<u>001</u>	Nov 26, 1997
-----------	--	------	-------------	----------------	------------	--------------

TRIDERM

<u>AT</u>		CROWN LABS	<u>0.025%</u>	<u>A088042</u>	<u>002</u>	Mar 25, 2015
-----------	--	------------	---------------	----------------	------------	--------------

<u>AT</u>			<u>0.1%</u>	<u>A088042</u>	<u>001</u>	Mar 19, 1984
-----------	--	--	-------------	----------------	------------	--------------

<u>AT</u>			<u>0.5%</u>	<u>A088042</u>	<u>003</u>	Mar 25, 2015
-----------	--	--	-------------	----------------	------------	--------------

FOR SUSPENSION, EXTENDED RELEASE; INTRA-ARTICULAR

ZILRETTA

+	PACIRA PHARMS INC	32MG/VIAL	N208845	001	Oct 06, 2017	
---	-------------------	-----------	---------	-----	--------------	--

INJECTABLE; INJECTION

KENALOG-40

<u>AB</u>	+	APOTHECON	<u>40MG/ML</u>	<u>N014901</u>	<u>001</u>	
-----------	---	-----------	----------------	----------------	------------	--

TRIAMCINOLONE ACETONIDE

<u>AB</u>		AMNEAL	<u>40MG/ML</u>	<u>A207550</u>	<u>001</u>	Dec 11, 2017
-----------	--	--------	----------------	----------------	------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

TRIAMCINOLONE ACETONIDE

INJECTABLE; INJECTION

TRIAMCINOLONE ACETONIDE

AB	EUGIA PHARMA	40MG/ML	A212400 001	Jul 05, 2022
AB	HIKMA	40MG/ML	A215254 001	Dec 09, 2024
AB	LONG GROVE PHARMS	40MG/ML	A213543 001	Jan 19, 2022
AB	MYLAN LABS LTD	40MG/ML	A212567 001	Nov 02, 2022
AB	TEVA PHARMS USA	40MG/ML	A209852 001	Oct 05, 2018
	KENALOG-10			
	+ APOTHECON	10MG/ML	N012041 001	
	KENALOG-80			
	+! APOTHECON	80MG/ML	N014901 002	Apr 12, 2019
	INJECTABLE; INTRAVITREAL			
	TRIESENCE			
	+! HARROW EYE	40MG/ML (40MG/ML)	N022048 001	Nov 29, 2007
	LOTION; TOPICAL			

TRIAMCINOLONE ACETONIDE

AT	COSETTE	0.1%	A089129 001	Aug 14, 1986
AT	EPIC PHARMA LLC	0.025%	A202374 001	May 08, 2013
AT		0.1%	A202374 002	May 08, 2013
AT	FOUGERA PHARMS	0.025%	A040467 001	Apr 21, 2003
AT		0.1%	A040467 002	Apr 21, 2003
AT	MICRO LABS	0.1%	A040672 002	Dec 13, 2006
AT	QUAGEN	0.025%	A213559 001	Jul 01, 2020
AT		0.1%	A213559 002	Jul 01, 2020
AT	! WOCKHARDT BIO AG	0.025%	A088450 001	Apr 01, 1985
AT	!	0.1%	A088451 001	Apr 03, 1985

OINTMENT; TOPICAL

TRIAMCINOLONE ACETONIDE

AT	CHARTWELL RX	0.05%	A214532 001	Jan 27, 2023
AT		0.1%	A207365 001	Oct 12, 2018
AT	CINTEX SVCS	0.05%	A213619 001	Nov 10, 2020
AT	COSETTE	0.025%	A089795 001	Dec 23, 1988
AT		0.1%	A089796 001	Dec 23, 1988
AT	! ENCUBE	0.05%	A212384 001	Nov 29, 2019
AT		0.1%	A205373 001	May 13, 2016
AT	FOUGERA PHARMS	0.025%	A085691 001	
AT		0.1%	A085691 003	
AT		0.5%	A085691 002	
AT	GLENMARK PHARMS LTD	0.5%	A206379 001	Jul 22, 2016
AT	GLENMARK SPECLT	0.1%	A208320 001	Aug 22, 2017
AT	MACLEODS PHARMS LTD	0.025%	A209828 001	Nov 23, 2018
AT		0.05%	A216625 001	Aug 18, 2023
AT		0.1%	A209828 002	Nov 23, 2018
AT		0.5%	A209828 003	Nov 23, 2018
AT	PADAGIS ISRAEL	0.05%	A212460 001	Feb 05, 2021
AT	+! PADAGIS US	0.025%	A087385 002	
AT	+!	0.1%	A087385 003	
AT	+!	0.5%	A087385 001	
AT	STRIDES PHARMA	0.05%	A212356 001	Jun 01, 2020
AT	TARO	0.1%	A040037 001	Sep 30, 1994

PASTE; DENTAL

TRIAMCINOLONE ACETONIDE

AT	COSETTE	0.1%	A205592 001	Jan 12, 2017
AT	QUAGEN	0.1%	A214582 001	Dec 09, 2022
AT	RISING	0.1%	A040771 001	Jul 01, 2010
AT	! TARO	0.1%	A070730 001	Oct 01, 1986

SPRAY; TOPICAL

TRIAMCINOLONE ACETONIDE

AT	! PADAGIS ISRAEL	0.147MG/GM	A205782 001	Apr 13, 2015
AT	RISING	0.147MG/GM	A206786 001	Sep 08, 2017
AT	SCIEGEN PHARMS INC	0.147MG/GM	A207094 001	Dec 07, 2016

SUSPENSION; INJECTION

XIPERE

	+! BAUSCH AND LOMB INC	40MG/ML	N211950 001	Oct 22, 2021
--	------------------------	---------	-------------	--------------

TRIAMTERENE

CAPSULE; ORAL

DYRENIUM

AB	+ ADVANZ PHARMA	50MG	N013174 001	
AB	+!	100MG	N013174 002	

PRESCRIPTION DRUG PRODUCT LIST

TRIAMTERENE

CAPSULE; ORAL

TRIAMTERENE

<u>AB</u>	AGNITIO	<u>50MG</u>	<u>A211581 001</u>	Aug 19, 2019
<u>AB</u>		<u>100MG</u>	<u>A211581 002</u>	Aug 19, 2019
<u>AB</u>	BIOCON GENERICS	<u>50MG</u>	<u>A214768 001</u>	Jul 06, 2022
<u>AB</u>		<u>100MG</u>	<u>A214768 002</u>	Jul 06, 2022

TRIAZOLAM

TABLET; ORAL

HALCION

<u>AB</u>	+	PFIZER	<u>0.125MG</u>	<u>N017892 003</u>	Apr 26, 1985
<u>AB</u>	+	!	<u>0.25MG</u>	<u>N017892 001</u>	Nov 15, 1982

TRIAZOLAM

<u>AB</u>		BRECKENRIDGE	<u>0.125MG</u>	<u>A216890 001</u>	Nov 08, 2022
<u>AB</u>			<u>0.25MG</u>	<u>A216890 002</u>	Nov 08, 2022
<u>AB</u>		NOVAST LABS	<u>0.125MG</u>	<u>A214219 001</u>	Oct 20, 2020
<u>AB</u>			<u>0.25MG</u>	<u>A214219 002</u>	Oct 20, 2020
<u>AB</u>		ZYDUS PHARMS	<u>0.125MG</u>	<u>A213003 001</u>	Dec 28, 2022
<u>AB</u>			<u>0.25MG</u>	<u>A213003 002</u>	Dec 28, 2022

TRICLABENDAZOLE

TABLET; ORAL

EGATEN

	+	!	NOVARTIS	250MG	N208711 001	Feb 13, 2019
--	---	---	----------	-------	-------------	--------------

TRIENTINE HYDROCHLORIDE

CAPSULE; ORAL

SYPRINE

<u>AB</u>	+	!	BAUSCH	<u>250MG</u>	<u>N019194 001</u>	Nov 08, 1985
-----------	---	---	--------	--------------	--------------------	--------------

TRIENTINE HYDROCHLORIDE

<u>AB</u>			DR REDDYS	<u>250MG</u>	<u>A211076 001</u>	Jul 03, 2019
<u>AB</u>			ECI PHARMS LLC	<u>250MG</u>	<u>A209945 001</u>	Aug 13, 2021
<u>AB</u>			ENDO OPERATIONS	<u>250MG</u>	<u>A210096 001</u>	Sep 25, 2019
<u>AB</u>			HETERO LABS LTD III	<u>250MG</u>	<u>A216356 001</u>	Jun 23, 2022
<u>AB</u>			MSN	<u>250MG</u>	<u>A211134 001</u>	May 22, 2019
<u>AB</u>			NAVINTA LLC	<u>250MG</u>	<u>A211251 001</u>	Jan 16, 2019
<u>AB</u>			RISING	<u>250MG</u>	<u>A212238 001</u>	Feb 20, 2020
<u>AB</u>			WATSON LABS TEVA	<u>250MG</u>	<u>A207567 001</u>	Feb 07, 2018
<u>AB</u>			ZYDUS PHARMS	<u>250MG</u>	<u>A211554 001</u>	Apr 26, 2019
			RISING	500MG	A212238 002	Sep 22, 2023

TRIENTINE TETRAHYDROCHLORIDE

TABLET; ORAL

CUVRIOR

	+	!	ORPHALAN	300MG	N215760 001	Apr 28, 2022
--	---	---	----------	-------	-------------	--------------

TRIFAROTENE

CREAM; TOPICAL

AKLIEF

	+	!	GALDERMA LABS LP	0.005%	N211527 001	Oct 04, 2019
--	---	---	------------------	--------	-------------	--------------

TRIFLUOPERAZINE HYDROCHLORIDE

TABLET; ORAL

TRIFLUOPERAZINE HYDROCHLORIDE

<u>AB</u>			MYLAN	<u>EQ 1MG BASE</u>	<u>A040209 001</u>	Jul 07, 1997
<u>AB</u>				<u>EQ 2MG BASE</u>	<u>A040209 002</u>	Jul 07, 1997
<u>AB</u>				<u>EQ 5MG BASE</u>	<u>A040209 003</u>	Jul 07, 1997
<u>AB</u>	!			<u>EQ 10MG BASE</u>	<u>A040209 004</u>	Jul 07, 1997
<u>AB</u>			SANDOZ	<u>EQ 1MG BASE</u>	<u>A085785 001</u>	
<u>AB</u>				<u>EQ 2MG BASE</u>	<u>A085786 001</u>	
<u>AB</u>				<u>EQ 5MG BASE</u>	<u>A085789 001</u>	
<u>AB</u>				<u>EQ 10MG BASE</u>	<u>A085788 001</u>	

TRIFLURIDINE

SOLUTION/DROPS; OPHTHALMIC

TRIFLURIDINE

<u>AT</u>			SANDOZ	<u>1%</u>	<u>A074311 001</u>	Oct 06, 1995
-----------	--	--	--------	-----------	--------------------	--------------

VIROPTIC

<u>AT</u>	+	!	MONARCH PHARMS	<u>1%</u>	<u>N018299 001</u>	
-----------	---	---	----------------	-----------	--------------------	--

PRESCRIPTION DRUG PRODUCT LIST

TRIEPTANOIN

LIQUID; ORAL

DOJOLVI

+	!	ULTRAGENYX PHARM INC	100% w/w	N213687	001	Jun 30, 2020
---	---	-------------------------	----------	---------	-----	--------------

TRIHXYPHENIDYL HYDROCHLORIDE

ELIXIR; ORAL

TRIHXYPHENIDYL HYDROCHLORIDE

AA		MIKART	2MG/5ML	A040251	001	Sep 27, 1999
AA	!	PHARM ASSOC	2MG/5ML	A040177	001	Apr 17, 1997

TABLET; ORAL

TRIHXYPHENIDYL HYDROCHLORIDE

AA		NATCO PHARMA LTD	2MG	A091630	001	Nov 17, 2010
AA			5MG	A091630	002	Nov 17, 2010
AA		NOVITIUM PHARMA	2MG	A040254	001	Dec 24, 1998
AA			5MG	A040254	002	Dec 24, 1998
AA	!	WATSON LABS	2MG	A084363	001	
AA	!		5MG	A084364	001	

TRILACICLIB DIHYDROCHLORIDE

POWDER; INTRAVENOUS

COSELA

+	!	PHARMACOSMOS	EQ 300MG BASE/VIAL	N214200	001	Feb 12, 2021
---	---	--------------	--------------------	---------	-----	--------------

TRIMETHOBENZAMIDE HYDROCHLORIDE

CAPSULE; ORAL

TRIMETHOBENZAMIDE HYDROCHLORIDE

AB	!	HERITAGE PHARMA AVET	300MG	A205950	001	Nov 21, 2023
AB		LUPIN	300MG	A076546	001	Aug 20, 2003

INJECTABLE; INJECTION

TIGAN

+	!	ENDO OPERATIONS	100MG/ML	N017530	001	
---	---	-----------------	----------	---------	-----	--

TRIMETHOPRIM

TABLET; ORAL

TRIMETHOPRIM

AB	+	!	DR REDDYS LABS SA	100MG	N018679	001	Jul 30, 1982
AB			NOVEL LABS INC	100MG	A091437	001	Jun 15, 2011
AB			NOVITIUM PHARMA	100MG	A216393	001	Oct 28, 2022
AB			WATSON LABS	100MG	A070049	001	Jun 06, 1985

TRIMIPRAMINE MALEATE

CAPSULE; ORAL

TRIMIPRAMINE MALEATE

AB		BRECKENRIDGE	EQ 25MG BASE	A208127	001	Apr 15, 2016
AB	!		EQ 50MG BASE	A208127	002	Apr 15, 2016
AB			EQ 100MG BASE	A208127	003	Apr 15, 2016
AB		ELITE LABS INC	EQ 25MG BASE	A077361	001	Aug 02, 2006
AB			EQ 50MG BASE	A077361	002	Aug 02, 2006
AB			EQ 100MG BASE	A077361	003	Aug 02, 2006

TRIPTORELIN PAMOATE

FOR SUSPENSION, EXTENDED RELEASE; INTRAMUSCULAR

TRIPTODUR KIT

+	!	AZURITY	EQ 22.5MG BASE/VIAL	N208956	001	Jun 29, 2017
---	---	---------	---------------------	---------	-----	--------------

INJECTABLE; INTRAMUSCULAR

TRELSTAR

+	!	VERITY	EQ 3.75MG BASE/VIAL	N020715	001	Jun 15, 2000
+	!		EQ 11.25MG BASE/VIAL	N021288	001	Jun 29, 2001
+	!		EQ 22.5MG BASE/VIAL	N022437	001	Mar 10, 2010

TROFINETIDE

SOLUTION; ORAL

DAYBUE

+	!	ACADIA PHARMS INC	200MG/ML	N217026	001	Mar 10, 2023
---	---	-------------------	----------	---------	-----	--------------

TROMETHAMINE

SOLUTION; INJECTION

THAM

AP	+	!	HOSPIRA	18GM/500ML (3.6GM/100ML)	N013025	002	
-----------	---	---	---------	---------------------------------	----------------	------------	--

TROMETHAMINE

AP		B BRAUN MEDICAL INC	18GM/500ML (3.6GM/100ML)	A211558	001	Dec 09, 2024
AP		MILLA PHARMS	18GM/500ML (3.6GM/100ML)	A213116	001	Dec 31, 2024

PRESCRIPTION DRUG PRODUCT LIST

TROPICAMIDE

SOLUTION/DROPS;OPHTHALMIC

MYDRIACYL

AT	+ !	ALCON LABS INC	1%	A084306	001	
AT	+ !	SANDOZ	0.5%	A084305	001	

TROPICACYL

AT		EPIC PHARMA LLC	0.5%	A040314	001	Sep 29, 2000
AT		RISING	1%	A040315	001	Sep 29, 2000

TROPICAMIDE

AT		BAUSCH AND LOMB	0.5%	A040067	001	Jul 27, 1994
AT			1%	A040064	001	Jul 27, 1994
AT		SOMERSET THERAPS LLC	1%	A207524	001	Dec 12, 2019

TROSPIMUM CHLORIDE

CAPSULE, EXTENDED RELEASE;ORAL

TROSPIMUM CHLORIDE

AB	!	ACTAVIS LABS FL INC	60MG	A091289	001	Oct 12, 2012
AB		GRANULES	60MG	A213185	001	Apr 23, 2020
AB		PADAGIS US	60MG	A201291	001	May 24, 2013
AB		UTOPIC PHARMS	60MG	A214760	001	Apr 30, 2021

TABLET;ORAL

TROSPIMUM CHLORIDE

AB		APOTEX	20MG	A091513	001	Dec 06, 2011
AB		CHARTWELL RX	20MG	A215781	001	Feb 16, 2022
AB	!	GLENMARK PHARMS LTD	20MG	A091575	001	Aug 13, 2010
AB		HERITAGE PHARMS INC	20MG	A204945	001	Aug 30, 2016
AB		INVAGEN PHARMS	20MG	A091688	001	Aug 23, 2016
AB		MACLEODS PHARMS LTD	20MG	A206472	001	Jun 18, 2024
AB		PADAGIS US	20MG	A091573	001	Nov 17, 2010

TROSPIMUM CHLORIDE; XANOMELINE TARTRATE

CAPSULE;ORAL

COBENFY

+	BRISTOL-MYERS	20MG;EQ 50MG BASE	N216158	001	Sep 26, 2024
+		20MG;EQ 100MG BASE	N216158	002	Sep 26, 2024
+ !		30MG;EQ 125MG BASE	N216158	003	Sep 26, 2024

TRYPAN BLUE

SOLUTION;OPHTHALMIC

VISIONBLUE

+ !	DORC	0.06%	N021670	001	Dec 16, 2004
------------	------	-------	---------	-----	--------------

TUCATINIB

TABLET;ORAL

TUKYSA

+	SEAGEN	50MG	N213411	001	Apr 17, 2020
+ !		150MG	N213411	002	Apr 17, 2020

UBROGEPANT

TABLET;ORAL

UBRELVY

+	ABBVIE	50MG	N211765	001	Dec 23, 2019
+ !		100MG	N211765	002	Dec 23, 2019

ULIPRISTAL ACETATE

TABLET;ORAL

ELLA

AB	+ !	LAB HRA PHARMA	30MG	N022474	001	Aug 13, 2010
-----------	------------	----------------	-------------	----------------	------------	--------------

LOGILIA

AB		TEVA PHARMS USA	30MG	A207952	001	Feb 13, 2017
-----------	--	-----------------	-------------	----------------	------------	--------------

UMECLIDINIUM BROMIDE

POWDER; INHALATION

INCRUSE ELLIPTA

+ !	GLAXO GRP ENGLAND	EQ 0.0625MG BASE/INH	N205382	001	Apr 30, 2014
------------	-------------------	----------------------	---------	-----	--------------

UMECLIDINIUM BROMIDE; VILANTEROL TRIFENATATE

POWDER; INHALATION

ANORO ELLIPTA

+ !	GLAXOSMITHKLINE	EQ 0.0625MG BASE/INH;EQ 0.025MG BASE/INH	N203975	001	Dec 18, 2013
------------	-----------------	---	---------	-----	--------------

PRESCRIPTION DRUG PRODUCT LIST

UPADACITINIB

SOLUTION;ORAL

RINVOQ LQ

+! ABBVIE 1MG/ML N218347 001 Apr 26, 2024

TABLET, EXTENDED RELEASE;ORAL

RINVOQ

+ ABBVIE 15MG N211675 001 Aug 16, 2019

+ 30MG N211675 002 Jan 14, 2022

+! 45MG N211675 003 Mar 16, 2022

URIDINE TRIACETATE

GRANULE;ORAL

VISTOGARD

+! BTG INTL 10GM/PACKET N208159 001 Dec 11, 2015

XURIDEN

+! BTG INTL 2GM/PACKET N208169 001 Sep 04, 2015

URSODIOL

CAPSULE;ORAL

ACTIGALL**AB** +! TEVA BRANDED PHARM **300MG** **N019594 002** Dec 31, 1987URSODIOL**AB** ABHAI LLC **300MG** **A210707 001** May 17, 2018**AB** AMNEAL PHARMS CO **300MG** **A211301 001** Oct 16, 2018**AB** AUROBINDO PHARMA **300MG** **A214849 001** Nov 28, 2022

LTD

AB CHARTWELL RX **300MG** **A213555 001** Aug 17, 2020**AB** EPIC PHARMA **300MG** **A075517 001** Mar 14, 2000**AB** HIBROW HLTHCARE **300MG** **A212452 001** Oct 30, 2019**AB** LANNETT CO INC **300MG** **A079082 001** Dec 15, 2008**AB** RISING **300MG** **A213200 001** Feb 12, 2020**AB** RK PHARMA **300MG** **A214329 001** Jul 28, 2021**AB** STRIDES PHARMA **300MG** **A210344 001** Jan 22, 2021**AB** ZYDUS LIFESCIENCES **300MG** **A214295 001** Oct 16, 2020

! LGM PHARMA 200MG A205789 001 May 08, 2020

! 400MG A205789 002 May 08, 2020

TABLET;ORAL

URSO 250**AB** + ALLERGAN **250MG** **N020675 001** Dec 10, 1997URSO FORTE**AB** +! ALLERGAN **500MG** **N020675 002** Jul 21, 2004URSODIOL**AB** ENDO OPERATIONS **250MG** **A202540 001** Feb 14, 2013**AB** **500MG** **A202540 002** Feb 14, 2013**AB** EPIC PHARMA LLC **250MG** **A214717 001** Aug 14, 2024**AB** **500MG** **A214717 002** Aug 14, 2024**AB** GLENMARK PHARMS LTD **250MG** **A090801 001** Jul 12, 2011**AB** **500MG** **A090801 002** Jul 12, 2011**AB** STRIDES PHARMA **250MG** **A213504 001** Aug 20, 2020**AB** **500MG** **A213504 002** Aug 20, 2020**AB** ZYDUS LIFESCIENCES **250MG** **A211145 001** Oct 30, 2018**AB** **500MG** **A211145 002** Oct 30, 2018VADADUSTAT

TABLET;ORAL

VAFSEO

+ AKEBIA 150MG N215192 001 Mar 27, 2024

+ 300MG N215192 002 Mar 27, 2024

+! 450MG N215192 003 Mar 27, 2024

VALACYCLOVIR HYDROCHLORIDE

TABLET;ORAL

VALACYCLOVIR HYDROCHLORIDE**AB** APOTEX **EQ 500MG BASE** **A090500 001** Apr 04, 2014**AB** **EQ 1GM BASE** **A090500 002** Apr 04, 2014**AB** AUROBINDO PHARMA **EQ 500MG BASE** **A090682 001** May 24, 2010**AB** **EQ 1GM BASE** **A090682 002** May 24, 2010**AB** CHARTWELL RX **EQ 500MG BASE** **A090216 001** May 24, 2010**AB** **EQ 1GM BASE** **A090216 002** May 24, 2010**AB** GRAVITI PHARMS **EQ 500MG BASE** **A079012 001** May 24, 2010**AB** **EQ 1GM BASE** **A079012 002** May 24, 2010**AB** HETERO LABS LTD V **EQ 500MG BASE** **A203047 001** Apr 08, 2015**AB** **EQ 1GM BASE** **A203047 002** Apr 08, 2015**AB** JUBILANT GENERICS **EQ 500MG BASE** **A201506 001** Apr 03, 2012**AB** **EQ 1GM BASE** **A201506 002** Apr 03, 2012

PRESCRIPTION DRUG PRODUCT LIST

VALACYCLOVIR HYDROCHLORIDE

TABLET; ORAL

VALACYCLOVIR HYDROCHLORIDE

<u>AB</u>	MYLAN PHARMS INC	<u>EQ 500MG BASE</u>	<u>A078518 001</u>	May 24, 2010
<u>AB</u>		<u>EQ 1GM BASE</u>	<u>A078518 002</u>	May 24, 2010
<u>AB</u>	SUN PHARM INDS LTD	<u>EQ 500MG BASE</u>	<u>A076588 001</u>	Jan 31, 2007
<u>AB</u>		<u>EQ 1GM BASE</u>	<u>A076588 002</u>	Jan 31, 2007
<u>AB</u>	YILING	<u>EQ 500MG BASE</u>	<u>A209553 001</u>	Mar 18, 2020
<u>AB</u>		<u>EQ 1GM BASE</u>	<u>A209553 002</u>	Mar 18, 2020
<u>AB</u>	ZYDUS LIFESCIENCES	<u>EQ 500MG BASE</u>	<u>A079137 001</u>	Dec 29, 2017
<u>AB</u>		<u>EQ 1GM BASE</u>	<u>A079137 002</u>	Dec 29, 2017

VALTREX

<u>AB</u>	+ GLAXOSMITHKLINE	<u>EQ 500MG BASE</u>	<u>N020487 001</u>	Jun 23, 1995
<u>AB</u>	+!	<u>EQ 1GM BASE</u>	<u>N020487 002</u>	Jun 23, 1995

VALBENAZINE TOSYLATE

CAPSULE; ORAL

INGREZZA

<u>AB</u>	+ NEUROCRINE	<u>EQ 40MG BASE</u>	<u>N209241 001</u>	Apr 11, 2017
<u>AB</u>	+	<u>EQ 60MG BASE</u>	<u>N209241 003</u>	Apr 23, 2021
<u>AB</u>	+!	<u>EQ 80MG BASE</u>	<u>N209241 002</u>	Oct 04, 2017

VALBENAZINE TOSYLATE

<u>AB</u>	LUPIN LTD	<u>EQ 40MG BASE</u>	<u>A216064 001</u>	Apr 05, 2024
<u>AB</u>		<u>EQ 80MG BASE</u>	<u>A216064 002</u>	Apr 05, 2024
<u>AB</u>	ZYDUS LIFESCIENCES	<u>EQ 40MG BASE</u>	<u>A216137 001</u>	Aug 07, 2024
<u>AB</u>		<u>EQ 60MG BASE</u>	<u>A216137 002</u>	Aug 07, 2024
<u>AB</u>		<u>EQ 80MG BASE</u>	<u>A216137 003</u>	Aug 07, 2024
	INGREZZA SPRINKLE			
	+ NEUROCRINE	EQ 40MG BASE	N218390 001	Apr 30, 2024
	+	EQ 60MG BASE	N218390 002	Apr 30, 2024
	+!	EQ 80MG BASE	N218390 003	Apr 30, 2024

VALGANCICLOVIR HYDROCHLORIDE

FOR SOLUTION; ORAL

VALCYTE

<u>AB</u>	+! CHEPLAPHARM	<u>50MG/ML</u>	<u>N022257 001</u>	Aug 28, 2009
-----------	----------------	----------------	--------------------	--------------

VALGANCICLOVIR HYDROCHLORIDE

<u>AB</u>	ACTAVIS LABS FL INC	<u>50MG/ML</u>	<u>A205220 001</u>	Jul 18, 2016
<u>AB</u>	AJANTA PHARMA LTD	<u>50MG/ML</u>	<u>A212890 001</u>	Jan 13, 2020
<u>AB</u>	APPCO	<u>50MG/ML</u>	<u>A216317 001</u>	Apr 11, 2023
<u>AB</u>	AUROBINDO PHARMA	<u>50MG/ML</u>	<u>A215124 001</u>	Nov 17, 2022
<u>AB</u>	GRANULES	<u>50MG/ML</u>	<u>A213306 001</u>	Jan 31, 2020
<u>AB</u>	HETERO LABS LTD V	<u>50MG/ML</u>	<u>A211475 001</u>	Oct 04, 2022
<u>AB</u>	MSN	<u>50MG/ML</u>	<u>A210169 001</u>	Feb 17, 2022

TABLET; ORAL

VALCYTE

<u>AB</u>	+! CHEPLAPHARM	<u>EQ 450MG BASE</u>	<u>N021304 001</u>	Mar 29, 2001
-----------	----------------	----------------------	--------------------	--------------

VALGANCICLOVIR HYDROCHLORIDE

<u>AB</u>	AJANTA PHARMA LTD	<u>EQ 450MG BASE</u>	<u>A212234 001</u>	Dec 26, 2019
<u>AB</u>	AUROBINDO PHARMA LTD	<u>EQ 450MG BASE</u>	<u>A204750 001</u>	Mar 31, 2016
<u>AB</u>	DR REDDYS	<u>EQ 450MG BASE</u>	<u>A203511 001</u>	Nov 04, 2014
<u>AB</u>		<u>EQ 450MG BASE</u>	<u>A206876 001</u>	Dec 12, 2017
<u>AB</u>	HETERO LABS LTD V	<u>EQ 450MG BASE</u>	<u>A205166 001</u>	Mar 18, 2016
<u>AB</u>	SOMERSET THERAPS LLC	<u>EQ 450MG BASE</u>	<u>A212876 001</u>	Jun 11, 2024
<u>AB</u>	STRIDES PHARMA	<u>EQ 450MG BASE</u>	<u>A200790 001</u>	Nov 04, 2014

VALPROATE SODIUM

INJECTABLE; INJECTION

VALPROATE SODIUM

<u>AP</u>	FRESENIUS KABI USA	<u>EQ 100MG BASE/ML</u>	<u>A076539 001</u>	Jun 26, 2003
<u>AP</u>	! HIKMA FARMACEUTICA	<u>EQ 100MG BASE/ML</u>	<u>A078523 001</u>	Feb 17, 2010
<u>AP</u>	SAGENT	<u>EQ 100MG BASE/ML</u>	<u>A076295 001</u>	Nov 14, 2002

VALPROIC ACID

CAPSULE; ORAL

VALPROIC ACID

<u>AB</u>	! BIONPHARMA	<u>250MG</u>	<u>A073484 001</u>	Jun 29, 1993
<u>AB</u>	CATALENT	<u>250MG</u>	<u>A073229 001</u>	Oct 29, 1991

SYRUP; ORAL

VALPROIC ACID

<u>AA</u>	ANI PHARMS	<u>250MG/5ML</u>	<u>A073178 001</u>	Aug 25, 1992
<u>AA</u>	CHARTWELL RX	<u>250MG/5ML</u>	<u>A075782 001</u>	Dec 22, 2000
<u>AA</u>	! PHARM ASSOC	<u>250MG/5ML</u>	<u>A075379 001</u>	Dec 15, 2000

PRESCRIPTION DRUG PRODUCT LIST

VALPROIC ACID

SYRUP; ORAL

VALPROIC ACID

<u>AA</u>	QUAGEN	<u>250MG/5ML</u>	<u>A090517 001</u>	May 28, 2010
-----------	--------	------------------	--------------------	--------------

VALRUBICIN

SOLUTION; INTRAVESICAL

VALRUBICIN

<u>AO</u>	HIKMA	<u>40MG/ML</u>	<u>A206430 001</u>	Apr 19, 2019
-----------	-------	----------------	--------------------	--------------

VALSTAR PRESERVATIVE FREE

<u>AO</u>	+! ENDO OPERATIONS	<u>40MG/ML</u>	<u>N020892 001</u>	Sep 25, 1998
-----------	--------------------	----------------	--------------------	--------------

VALSARTAN

SOLUTION; ORAL

VALSARTAN

! NOVITIUM PHARMA

20MG/5ML

A214102 001 Nov 02, 2021

TABLET; ORAL

DIOVAN

<u>AB</u>	+ NOVARTIS	<u>40MG</u>	<u>N021283 004</u>	Aug 14, 2002
-----------	------------	-------------	--------------------	--------------

<u>AB</u>	+	<u>80MG</u>	<u>N021283 001</u>	Jul 18, 2001
-----------	---	-------------	--------------------	--------------

<u>AB</u>	+	<u>160MG</u>	<u>N021283 002</u>	Jul 18, 2001
-----------	---	--------------	--------------------	--------------

<u>AB</u>	+!	<u>320MG</u>	<u>N021283 003</u>	Jul 18, 2001
-----------	----	--------------	--------------------	--------------

VALSARTAN

<u>AB</u>	ALEMBIC	<u>40MG</u>	<u>A091367 001</u>	Jan 05, 2015
-----------	---------	-------------	--------------------	--------------

<u>AB</u>		<u>80MG</u>	<u>A091367 002</u>	Jan 05, 2015
-----------	--	-------------	--------------------	--------------

<u>AB</u>		<u>160MG</u>	<u>A091367 003</u>	Jan 05, 2015
-----------	--	--------------	--------------------	--------------

<u>AB</u>		<u>320MG</u>	<u>A091367 004</u>	Jan 05, 2015
-----------	--	--------------	--------------------	--------------

<u>AB</u>	ALKEM LABS LTD	<u>40MG</u>	<u>A205536 001</u>	Mar 12, 2019
-----------	----------------	-------------	--------------------	--------------

<u>AB</u>		<u>80MG</u>	<u>A205536 002</u>	Mar 12, 2019
-----------	--	-------------	--------------------	--------------

<u>AB</u>		<u>160MG</u>	<u>A205536 003</u>	Mar 12, 2019
-----------	--	--------------	--------------------	--------------

<u>AB</u>		<u>320MG</u>	<u>A205536 004</u>	Mar 12, 2019
-----------	--	--------------	--------------------	--------------

<u>AB</u>	AMNEAL PHARMS	<u>40MG</u>	<u>A204011 001</u>	Jan 11, 2016
-----------	---------------	-------------	--------------------	--------------

<u>AB</u>		<u>80MG</u>	<u>A204011 002</u>	Jan 11, 2016
-----------	--	-------------	--------------------	--------------

<u>AB</u>		<u>160MG</u>	<u>A204011 003</u>	Jan 11, 2016
-----------	--	--------------	--------------------	--------------

<u>AB</u>		<u>320MG</u>	<u>A204011 004</u>	Jan 11, 2016
-----------	--	--------------	--------------------	--------------

<u>AB</u>	AUROBINDO PHARMA LTD	<u>40MG</u>	<u>A202223 001</u>	Jan 05, 2015
-----------	----------------------	-------------	--------------------	--------------

<u>AB</u>		<u>80MG</u>	<u>A202223 002</u>	Jan 05, 2015
-----------	--	-------------	--------------------	--------------

<u>AB</u>		<u>160MG</u>	<u>A202223 003</u>	Jan 05, 2015
-----------	--	--------------	--------------------	--------------

<u>AB</u>		<u>320MG</u>	<u>A202223 004</u>	Jan 05, 2015
-----------	--	--------------	--------------------	--------------

<u>AB</u>	DR REDDYS	<u>40MG</u>	<u>A201618 001</u>	Oct 05, 2021
-----------	-----------	-------------	--------------------	--------------

<u>AB</u>		<u>80MG</u>	<u>A201618 002</u>	Oct 05, 2021
-----------	--	-------------	--------------------	--------------

<u>AB</u>		<u>160MG</u>	<u>A201618 003</u>	Oct 05, 2021
-----------	--	--------------	--------------------	--------------

<u>AB</u>		<u>320MG</u>	<u>A201618 004</u>	Oct 05, 2021
-----------	--	--------------	--------------------	--------------

<u>AB</u>	HETERO LABS LTD V	<u>40MG</u>	<u>A203311 001</u>	Jan 05, 2015
-----------	-------------------	-------------	--------------------	--------------

<u>AB</u>		<u>80MG</u>	<u>A203311 002</u>	Jan 05, 2015
-----------	--	-------------	--------------------	--------------

<u>AB</u>		<u>160MG</u>	<u>A203311 003</u>	Jan 05, 2015
-----------	--	--------------	--------------------	--------------

<u>AB</u>		<u>320MG</u>	<u>A203311 004</u>	Jan 05, 2015
-----------	--	--------------	--------------------	--------------

<u>AB</u>	JUBILANT GENERICS	<u>40MG</u>	<u>A203536 001</u>	Jan 05, 2015
-----------	-------------------	-------------	--------------------	--------------

<u>AB</u>		<u>80MG</u>	<u>A203536 002</u>	Jan 05, 2015
-----------	--	-------------	--------------------	--------------

<u>AB</u>		<u>160MG</u>	<u>A203536 003</u>	Jan 05, 2015
-----------	--	--------------	--------------------	--------------

<u>AB</u>		<u>320MG</u>	<u>A203536 004</u>	Jan 05, 2015
-----------	--	--------------	--------------------	--------------

<u>AB</u>	LUPIN LTD	<u>40MG</u>	<u>A201677 001</u>	Jan 05, 2015
-----------	-----------	-------------	--------------------	--------------

<u>AB</u>		<u>80MG</u>	<u>A201677 002</u>	Jan 05, 2015
-----------	--	-------------	--------------------	--------------

<u>AB</u>		<u>160MG</u>	<u>A201677 003</u>	Jan 05, 2015
-----------	--	--------------	--------------------	--------------

<u>AB</u>		<u>320MG</u>	<u>A201677 004</u>	Jan 05, 2015
-----------	--	--------------	--------------------	--------------

<u>AB</u>	MACLEODS PHARMS LTD	<u>40MG</u>	<u>A202696 001</u>	Sep 16, 2016
-----------	---------------------	-------------	--------------------	--------------

<u>AB</u>		<u>80MG</u>	<u>A202696 002</u>	Sep 16, 2016
-----------	--	-------------	--------------------	--------------

<u>AB</u>		<u>160MG</u>	<u>A202696 003</u>	Sep 16, 2016
-----------	--	--------------	--------------------	--------------

<u>AB</u>		<u>320MG</u>	<u>A202696 004</u>	Sep 16, 2016
-----------	--	--------------	--------------------	--------------

<u>AB</u>	MYLAN	<u>40MG</u>	<u>A090866 001</u>	Jan 05, 2015
-----------	-------	-------------	--------------------	--------------

<u>AB</u>		<u>80MG</u>	<u>A090866 002</u>	Jan 05, 2015
-----------	--	-------------	--------------------	--------------

<u>AB</u>		<u>160MG</u>	<u>A090866 003</u>	Jan 05, 2015
-----------	--	--------------	--------------------	--------------

<u>AB</u>		<u>320MG</u>	<u>A090866 004</u>	Jan 05, 2015
-----------	--	--------------	--------------------	--------------

<u>AB</u>	OHM LABS INC	<u>40MG</u>	<u>A077492 001</u>	Jun 26, 2014
-----------	--------------	-------------	--------------------	--------------

<u>AB</u>		<u>80MG</u>	<u>A077492 002</u>	Jun 26, 2014
-----------	--	-------------	--------------------	--------------

<u>AB</u>		<u>160MG</u>	<u>A077492 003</u>	Jun 26, 2014
-----------	--	--------------	--------------------	--------------

<u>AB</u>		<u>320MG</u>	<u>A077492 004</u>	Jun 26, 2014
-----------	--	--------------	--------------------	--------------

<u>AB</u>	PRINSTON INC	<u>40MG</u>	<u>A204821 001</u>	Jun 09, 2015
-----------	--------------	-------------	--------------------	--------------

<u>AB</u>		<u>80MG</u>	<u>A204821 002</u>	Jun 09, 2015
-----------	--	-------------	--------------------	--------------

<u>AB</u>		<u>160MG</u>	<u>A204821 003</u>	Jun 09, 2015
-----------	--	--------------	--------------------	--------------

<u>AB</u>		<u>320MG</u>	<u>A204821 004</u>	Jun 09, 2015
-----------	--	--------------	--------------------	--------------

<u>AB</u>	RYAN LABS	<u>40MG</u>	<u>A218169 001</u>	Aug 05, 2024
-----------	-----------	-------------	--------------------	--------------

<u>AB</u>		<u>80MG</u>	<u>A218169 002</u>	Aug 05, 2024
-----------	--	-------------	--------------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

VALSARTAN

TABLET; ORAL

VALSARTAN

<u>AB</u>		<u>160MG</u>	<u>A218169 003</u>	Aug 05, 2024
<u>AB</u>		<u>320MG</u>	<u>A218169 004</u>	Aug 05, 2024
<u>AB</u>	SCIEGEN PHARMS INC	<u>40MG</u>	<u>A204038 001</u>	Oct 27, 2021
<u>AB</u>		<u>80MG</u>	<u>A204038 002</u>	Oct 27, 2021
<u>AB</u>		<u>160MG</u>	<u>A204038 003</u>	Oct 27, 2021
<u>AB</u>		<u>320MG</u>	<u>A204038 004</u>	Oct 27, 2021
<u>AB</u>	SQUARE PHARMS	<u>40MG</u>	<u>A205347 001</u>	Apr 09, 2018
<u>AB</u>		<u>80MG</u>	<u>A205347 002</u>	Apr 09, 2018
<u>AB</u>		<u>160MG</u>	<u>A205347 003</u>	Apr 09, 2018
<u>AB</u>		<u>320MG</u>	<u>A205347 004</u>	Apr 09, 2018
<u>AB</u>	ZYDUS LIFESCIENCES	<u>40MG</u>	<u>A218991 001</u>	Jul 22, 2024
<u>AB</u>		<u>80MG</u>	<u>A218991 002</u>	Jul 22, 2024
<u>AB</u>		<u>160MG</u>	<u>A218991 003</u>	Jul 22, 2024
<u>AB</u>		<u>320MG</u>	<u>A218991 004</u>	Jul 22, 2024

VAMOROLONE

SUSPENSION; ORAL

AGAMREE

+! CATALYST PHARMS 40MG/ML N215239 001 Oct 26, 2023

VANCOMYCIN HYDROCHLORIDE

CAPSULE; ORAL

VANCOICIN HYDROCHLORIDE

<u>AB</u>	+ ANI PHARMS	<u>EQ 125MG BASE</u>	<u>N050606 001</u>	Apr 15, 1986
<u>AB</u>	+!	<u>EQ 250MG BASE</u>	<u>N050606 002</u>	Apr 15, 1986

VANCOMYCIN HYDROCHLORIDE

<u>AB</u>	LUPIN LTD	<u>EQ 125MG BASE</u>	<u>A090439 001</u>	Jan 28, 2015
<u>AB</u>		<u>EQ 250MG BASE</u>	<u>A090439 002</u>	Jan 28, 2015
<u>AB</u>	ORIENT PHARMA CO LTD	<u>EQ 125MG BASE</u>	<u>A210729 001</u>	Apr 29, 2019
<u>AB</u>		<u>EQ 250MG BASE</u>	<u>A210729 002</u>	Apr 29, 2019
<u>AB</u>	STRIDES PHARMA	<u>EQ 125MG BASE</u>	<u>A065490 001</u>	Apr 09, 2012
<u>AB</u>		<u>EQ 250MG BASE</u>	<u>A065490 002</u>	Apr 09, 2012

FOR SOLUTION; ORAL

VANCOICIN HYDROCHLORIDE

<u>AA</u>	! ANI PHARMS	<u>EQ 250MG BASE/5ML</u>	<u>A061667 002</u>	Jul 13, 1983
-----------	--------------	--------------------------	--------------------	--------------

VANCOMYCIN HYDROCHLORIDE

<u>AA</u>	AMNEAL	<u>EQ 250MG BASE/5ML</u>	<u>A215338 001</u>	Jun 23, 2023
-----------	--------	--------------------------	--------------------	--------------

FIRVANO KIT

<u>AB</u>	+! AZURITY	<u>EQ 25MG BASE/ML</u>	<u>N208910 001</u>	Jan 26, 2018
<u>AB</u>	+!	<u>EQ 50MG BASE/ML</u>	<u>N208910 002</u>	Jan 26, 2018

VANCOMYCIN HYDROCHLORIDE

<u>AB</u>	ALKEM LABS LTD	<u>EQ 25MG BASE/ML</u>	<u>A214913 001</u>	Nov 14, 2022
<u>AB</u>		<u>EQ 50MG BASE/ML</u>	<u>A214913 002</u>	Nov 14, 2022

INJECTABLE; INJECTION

VANCOMYCIN HYDROCHLORIDE

<u>AP</u>	ASPIRO	<u>EQ 500MG BASE/VIAL</u>	<u>A216591 001</u>	Jul 06, 2022
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>A216591 002</u>	Jul 06, 2022
<u>AP</u>	EUGIA PHARMA	<u>EQ 500MG BASE/VIAL</u>	<u>A205780 001</u>	Mar 31, 2016
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>A205780 002</u>	Mar 31, 2016
<u>AP</u>		<u>EQ 5GM BASE/VIAL</u>	<u>A205779 001</u>	Mar 29, 2016
<u>AP</u>		<u>EQ 10GM BASE/VIAL</u>	<u>A205779 002</u>	Mar 29, 2016
<u>AP</u>	! FRESENIUS KABI USA	<u>EQ 500MG BASE/VIAL</u>	<u>A062663 001</u>	Mar 17, 1987
<u>AP</u>		<u>EQ 750MG BASE/VIAL</u>	<u>A062663 005</u>	Aug 17, 2016
<u>AP</u>	!	<u>EQ 1GM BASE/VIAL</u>	<u>A062663 002</u>	Jul 31, 1987
<u>AP</u>	!	<u>EQ 5GM BASE/VIAL</u>	<u>A062663 003</u>	Jun 03, 1988
<u>AP</u>	!	<u>EQ 10GM BASE/VIAL</u>	<u>A062663 004</u>	Nov 28, 1997
<u>AP</u>	GLAND PHARMA LTD	<u>EQ 500MG BASE/VIAL</u>	<u>A205694 001</u>	Jan 21, 2016
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>A205694 002</u>	Jan 21, 2016
<u>AP</u>		<u>EQ 5GM BASE/VIAL</u>	<u>A205694 003</u>	Jun 06, 2023
<u>AP</u>		<u>EQ 10GM BASE/VIAL</u>	<u>A205694 004</u>	Jun 06, 2023
<u>AP</u>	HIKMA	<u>EQ 500MG BASE/VIAL</u>	<u>A204107 001</u>	Dec 28, 2015
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>A203300 002</u>	Aug 11, 2020
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>A204107 002</u>	Dec 28, 2015
<u>AP</u>		<u>EQ 5GM BASE/VIAL</u>	<u>A204360 001</u>	Oct 15, 2018
<u>AP</u>		<u>EQ 10GM BASE/VIAL</u>	<u>A204360 002</u>	Oct 15, 2018
<u>AP</u>	HIKMA PHARMS	<u>EQ 750MG BASE/VIAL</u>	<u>A206616 001</u>	Oct 03, 2018
<u>AP</u>	! HOSPIRA	<u>EQ 500MG BASE/VIAL</u>	<u>A062911 001</u>	Aug 04, 1988
<u>AP</u>	!	<u>EQ 500MG BASE/VIAL</u>	<u>A062931 001</u>	Oct 29, 1992
<u>AP</u>		<u>EQ 750MG BASE/VIAL</u>	<u>A062912 002</u>	Jan 07, 2009
<u>AP</u>		<u>EQ 750MG BASE/VIAL</u>	<u>A062933 002</u>	May 27, 2009

PRESCRIPTION DRUG PRODUCT LIST

VANCOMYCIN HYDROCHLORIDE

INJECTABLE; INJECTION

VANCOMYCIN HYDROCHLORIDE

<u>AP</u>	!		<u>EQ 1GM BASE/VIAL</u>	<u>A062912 001</u>	Aug 04, 1988
<u>AP</u>	!		<u>EQ 1GM BASE/VIAL</u>	<u>A062933 001</u>	Oct 29, 1992
<u>AP</u>	!		<u>EQ 5GM BASE/VIAL</u>	<u>A063076 001</u>	Dec 21, 1990
<u>AP</u>			<u>EQ 10GM BASE/VIAL</u>	<u>A063076 002</u>	Sep 14, 2020
<u>AP</u>		HOSPIRA INC	<u>EQ 10GM BASE/VIAL</u>	<u>A065455 001</u>	Apr 29, 2009
<u>AP</u>		MEITHEAL	<u>EQ 500MG BASE/VIAL</u>	<u>A215197 001</u>	Jul 28, 2022
<u>AP</u>			<u>EQ 750MG BASE/VIAL</u>	<u>A215195 001</u>	Sep 15, 2022
<u>AP</u>			<u>EQ 1GM BASE/VIAL</u>	<u>A215197 002</u>	Jul 28, 2022
<u>AP</u>			<u>EQ 5GM BASE/VIAL</u>	<u>A215196 001</u>	Jul 27, 2022
<u>AP</u>			<u>EQ 10GM BASE/VIAL</u>	<u>A215196 002</u>	Jul 27, 2022
<u>AP</u>		MYLAN LABS LTD	<u>EQ 500MG BASE/VIAL</u>	<u>A065397 001</u>	Dec 30, 2008
<u>AP</u>			<u>EQ 1GM BASE/VIAL</u>	<u>A065397 002</u>	Dec 30, 2008
<u>AP</u>			<u>EQ 5GM BASE/VIAL</u>	<u>A065432 001</u>	Dec 30, 2008
<u>AP</u>	!		<u>EQ 10GM BASE/VIAL</u>	<u>A091554 001</u>	Sep 19, 2011
<u>AP</u>		SAGENT PHARMS	<u>EQ 5GM BASE/VIAL</u>	<u>A200837 001</u>	Aug 10, 2012
<u>AP</u>			<u>EQ 10GM BASE/VIAL</u>	<u>A200837 002</u>	Sep 02, 2014
<u>AP</u>		SLATE RUN PHARMA	<u>EQ 500MG BASE/VIAL</u>	<u>A212332 001</u>	Jun 12, 2019
<u>AP</u>			<u>EQ 1GM BASE/VIAL</u>	<u>A212332 002</u>	Jun 12, 2019
<u>AP</u>			<u>EQ 5GM BASE/VIAL</u>	<u>A215821 001</u>	Nov 18, 2021
<u>AP</u>			<u>EQ 10GM BASE/VIAL</u>	<u>A215821 002</u>	Nov 18, 2021
<u>AP</u>		XELLIA PHARMS APS	<u>EQ 5GM BASE/VIAL</u>	<u>A204125 001</u>	Dec 28, 2015
<u>AP</u>			<u>EQ 10GM BASE/VIAL</u>	<u>A204125 002</u>	Dec 28, 2015
	!	HOSPIRA	EQ 1.5GM BASE/VIAL	A062912 003	Jul 10, 2020

POWDER; INTRAVENOUS

VANCOMYCIN HYDROCHLORIDE

<u>AP</u>		EUGIA PHARMA	<u>EQ 1.25GM BASE/VIAL</u>	<u>A217401 001</u>	Aug 04, 2023
<u>AP</u>			<u>EQ 1.5GM BASE/VIAL</u>	<u>A217401 002</u>	Aug 04, 2023
<u>AP</u>		HIKMA	<u>EQ 1.25GM BASE/VIAL</u>	<u>A217489 001</u>	Jun 29, 2023
<u>AP</u>			<u>EQ 1.5GM BASE/VIAL</u>	<u>A217489 002</u>	Jun 29, 2023
<u>AP</u>	+	MYLAN LABS LTD	<u>EQ 1.25GM BASE/VIAL</u>	<u>N209481 003</u>	Jul 10, 2018
<u>AP</u>	+		<u>EQ 1.5GM BASE/VIAL</u>	<u>N209481 004</u>	Jul 10, 2018
	+		EQ 750MG BASE/VIAL	N209481 002	Jul 10, 2018
	+		EQ 1.75GM BASE/VIAL	N209481 005	Jun 26, 2024
	+		EQ 2GM BASE/VIAL	N209481 006	Jun 26, 2024
		VANCOMYCIN HYDROCHLORIDE IN PLASTIC CONTAINER			
		SAMSON MEDCL	EQ 100GM BASE	A091532 001	Jan 06, 2016
		POWDER; INTRAVENOUS, ORAL			
		VANCOMYCIN HYDROCHLORIDE			
	+	ZHEJIANG NOVUS PHARM	EQ 500MG BASE/VIAL	N210274 001	Jan 20, 2023
	+		EQ 1GM BASE/VIAL	N210274 002	Jan 20, 2023
	+		EQ 5GM BASE/VIAL	N210274 003	Jan 20, 2023
	+		EQ 10GM BASE/VIAL	N210274 004	Jan 20, 2023

SOLUTION; INTRAVENOUS

VANCOMYCIN HYDROCHLORIDE

	+	HIKMA	EQ 500MG BASE/100ML (EQ 5MG BASE/ML)	N211962 001	Feb 15, 2019
	+		EQ 750MG BASE/150ML (EQ 5MG BASE/ML)	N211962 005	May 13, 2020
	+		EQ 1GM BASE/200ML (EQ 5MG BASE/ML)	N211962 002	Feb 15, 2019
	+		EQ 1.25GM BASE/250ML (EQ 5MG BASE/ML)	N211962 006	May 13, 2020
	+		EQ 1.5GM BASE/300ML (EQ 5MG BASE/ML)	N211962 003	Feb 15, 2019
	+		EQ 1.75GM BASE/350ML (EQ 5MG BASE/ML)	N211962 007	May 13, 2020
	+		EQ 2GM BASE/400ML (EQ 5MG BASE/ML)	N211962 004	Feb 15, 2019

VANCOMYCIN HYDROCHLORIDE IN PLASTIC CONTAINER

	+	BAXTER HLTHCARE	EQ 500MG BASE/100ML (EQ 5MG BASE/ML)	N050671 001	Apr 29, 1993
	+		EQ 750MG BASE/150ML (EQ 5MG BASE/ML)	N050671 002	Dec 20, 2010
	+		EQ 1GM BASE/200ML (EQ 5MG BASE/ML)	N050671 003	Mar 01, 1999
	+		EQ 1.25GM BASE/250ML (EQ 5MG BASE/ML)	N050671 004	Jan 25, 2024
	+		EQ 1.5GM BASE/300ML (EQ 5MG BASE/ML)	N050671 005	Jan 25, 2024

VANDETANIB

TABLET; ORAL

CAPRELSA

	+	GENZYME CORP	100MG	N022405 001	Apr 06, 2011
	+		300MG	N022405 002	Apr 06, 2011

PRESCRIPTION DRUG PRODUCT LIST

VARDENAFIL HYDROCHLORIDE

TABLET;ORAL

VARDENAFIL HYDROCHLORIDE

<u>AB</u>	ALEMBIC	<u>EQ 2.5MG BASE</u>	<u>A214031 001</u>	Aug 04, 2020
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A214031 002</u>	Aug 04, 2020
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A214031 003</u>	Aug 04, 2020
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A214031 004</u>	Aug 04, 2020
<u>AB</u>	CROSSMEDIKA SA	<u>EQ 2.5MG BASE</u>	<u>A209057 001</u>	Oct 31, 2018
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A209057 002</u>	Oct 31, 2018
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A209057 003</u>	Oct 31, 2018
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A209057 004</u>	Oct 31, 2018
<u>AB</u>	MACLEODS PHARMS LTD	<u>EQ 2.5MG BASE</u>	<u>A204632 001</u>	Oct 22, 2019
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A204632 002</u>	Oct 22, 2019
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A204632 003</u>	Oct 22, 2019
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A204632 004</u>	Oct 22, 2019
<u>AB</u>	TEVA PHARMS	<u>EQ 2.5MG BASE</u>	<u>A091347 001</u>	May 03, 2012
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A091347 002</u>	May 03, 2012
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A091347 003</u>	May 03, 2012
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A091347 004</u>	May 03, 2012
<u>AB</u>	ZYDUS PHARMS	<u>EQ 2.5MG BASE</u>	<u>A208960 001</u>	Oct 31, 2018
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A208960 002</u>	Oct 31, 2018
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A208960 003</u>	Oct 31, 2018
<u>AB</u>	!	<u>EQ 20MG BASE</u>	<u>A208960 004</u>	Oct 31, 2018

TABLET, ORALLY DISINTEGRATING;ORAL

VARDENAFIL HYDROCHLORIDE

<u>AB</u>	!	ALEMBIC	<u>10MG</u>	<u>A208324 001</u>	Nov 16, 2018
<u>AB</u>		MACLEODS PHARMS LTD	<u>10MG</u>	<u>A205988 001</u>	Mar 10, 2020

VARENICLINE TARTRATE

SPRAY;NASAL

TYRVAYA

+! OYSTER POINT PHARMA EQ 0.03MG BASE/SPRAY

N213978 001 Oct 15, 2021

TABLET;ORAL

VARENICLINE TARTRATE

<u>AB</u>	AJANTA PHARMA LTD	<u>EQ 0.5MG BASE</u>	<u>A213019 001</u>	Mar 19, 2024
<u>AB</u>		<u>EQ 1MG BASE</u>	<u>A213019 002</u>	Mar 19, 2024
<u>AB</u>	ALKEM LABS LTD	<u>EQ 0.5MG BASE</u>	<u>A214557 001</u>	Aug 23, 2023
<u>AB</u>		<u>EQ 1MG BASE</u>	<u>A214557 002</u>	Aug 23, 2023
<u>AB</u>	APOTEX	<u>EQ 0.5MG BASE</u>	<u>A201962 001</u>	Jan 25, 2023
<u>AB</u>		<u>EQ 1MG BASE</u>	<u>A201962 002</u>	Jan 25, 2023
<u>AB</u>	DR REDDYS	<u>EQ 0.5MG BASE</u>	<u>A215931 001</u>	Oct 01, 2024
<u>AB</u>		<u>EQ 1MG BASE</u>	<u>A215931 002</u>	Oct 01, 2024
<u>AB</u>	ENDO OPERATIONS	<u>EQ 0.5MG BASE</u>	<u>A201785 001</u>	Aug 11, 2021
<u>AB</u>	!	<u>EQ 1MG BASE</u>	<u>A201785 002</u>	Aug 11, 2021
<u>AB</u>	HETERO LABS LTD III	<u>EQ 0.5MG BASE</u>	<u>A214571 001</u>	Oct 23, 2023
<u>AB</u>		<u>EQ 1MG BASE</u>	<u>A214571 002</u>	Oct 23, 2023
<u>AB</u>	INDOCO	<u>EQ 0.5MG BASE</u>	<u>A219106 001</u>	Oct 29, 2024
<u>AB</u>		<u>EQ 1MG BASE</u>	<u>A219106 002</u>	Oct 29, 2024
<u>AB</u>	LUPIN LTD	<u>EQ 0.5MG BASE</u>	<u>A211862 001</u>	Dec 04, 2023
<u>AB</u>		<u>EQ 1MG BASE</u>	<u>A211862 002</u>	Dec 04, 2023
<u>AB</u>	MACLEODS PHARMS LTD	<u>EQ 0.5MG BASE</u>	<u>A215048 001</u>	Dec 09, 2024
<u>AB</u>		<u>EQ 1MG BASE</u>	<u>A215048 002</u>	Dec 09, 2024
<u>AB</u>	MANKIND PHARMA	<u>EQ 0.5MG BASE</u>	<u>A214255 001</u>	Aug 01, 2023
<u>AB</u>		<u>EQ 1MG BASE</u>	<u>A214255 002</u>	Aug 01, 2023
<u>AB</u>	MEDLEY PHARMS	<u>EQ 0.5MG BASE</u>	<u>A217151 001</u>	Jul 25, 2023
<u>AB</u>		<u>EQ 1MG BASE</u>	<u>A217151 002</u>	Jul 25, 2023
<u>AB</u>	MYLAN	<u>EQ 0.5MG BASE</u>	<u>A202019 001</u>	Feb 28, 2024
<u>AB</u>		<u>EQ 1MG BASE</u>	<u>A202019 002</u>	Feb 28, 2024
<u>AB</u>	NE RX PHARMA	<u>EQ 0.5MG BASE</u>	<u>A217283 001</u>	Mar 06, 2024
<u>AB</u>		<u>EQ 1MG BASE</u>	<u>A217283 002</u>	Mar 06, 2024
<u>AB</u>	PIRAMAL	<u>EQ 0.5MG BASE</u>	<u>A217115 001</u>	Jul 23, 2024
<u>AB</u>		<u>EQ 1MG BASE</u>	<u>A217115 002</u>	Jul 23, 2024
<u>AB</u>	ZYDUS	<u>EQ 0.5MG BASE</u>	<u>A216723 001</u>	Jun 12, 2023
<u>AB</u>		<u>EQ 1MG BASE</u>	<u>A216723 002</u>	Jun 12, 2023

VASOPRESSIN

SOLUTION;INTRAVENOUS

VASOPRESSIN

<u>AP</u>	AMNEAL	<u>20UNITS/ML (20UNITS/ML)</u>	<u>A212944 001</u>	Aug 05, 2022
<u>AP</u>	AMPHASTAR PHARMS INC	<u>20UNITS/ML (20UNITS/ML)</u>	<u>A211857 001</u>	Jul 18, 2022
<u>AP</u>	CIPLA	<u>20UNITS/100ML (0.2UNITS/ML)</u>	<u>A217987 001</u>	Dec 06, 2023
<u>AP</u>	DR REDDYS	<u>20UNITS/ML (20UNITS/ML)</u>	<u>A213988 001</u>	May 08, 2024
<u>AP</u>	EAGLE PHARMS	<u>20UNITS/ML (20UNITS/ML)</u>	<u>A211538 001</u>	Dec 15, 2021

PRESCRIPTION DRUG PRODUCT LIST

VASOPRESSIN

SOLUTION; INTRAVENOUS

VASOPRESSIN

<u>AP</u>	EUGIA PHARMA	<u>20UNITS/ML (20UNITS/ML)</u>	<u>A214314 001</u>	Aug 15, 2022
<u>AP</u>	FRESENIUS KABI USA	<u>20UNITS/ML (20UNITS/ML)</u>	<u>A213206 001</u>	May 26, 2023
<u>AP</u>	GLAND PHARMA LTD	<u>20UNITS/ML (20UNITS/ML)</u>	<u>A216963 001</u>	Feb 09, 2024

VASOSTRICT

<u>AP</u>	+! ENDO OPERATIONS	<u>20UNITS/ML (20UNITS/ML)</u>	<u>N204485 001</u>	Apr 17, 2014
<u>AP</u>	+!	<u>20UNITS/100ML (0.2UNITS/ML)</u>	<u>N204485 005</u>	Apr 21, 2021

VASOPRESSIN

+!	AM REGENT	20UNITS/ML (20UNITS/ML)	N212593 001	Aug 03, 2020
+!		200UNITS/10ML (20UNITS/ML)	N212593 002	Jun 09, 2023

VASOPRESSIN IN SODIUM CHLORIDE 0.9%

+!	BAXTER HLTHCARE CORP	20UNITS/100ML (0.2UNITS/ML)	N217569 001	Sep 29, 2023
+!		40UNITS/100ML (0.4UNITS/ML)	N217569 002	Sep 29, 2023
+!	LONG GROVE PHARMS	20UNITS/100ML (0.2UNITS/ML)	N217766 001	Jul 11, 2024
+!		40UNITS/100ML (0.4UNITS/ML)	N217766 002	Jul 11, 2024
+!		50UNITS/50ML (1UNIT/ML)	N217766 003	Jul 11, 2024

VASOSTRICT

+!	ENDO OPERATIONS	40UNITS/100ML (0.4UNITS/ML)	N204485 003	Apr 15, 2020
+!		200UNITS/10ML (20UNITS/ML)	N204485 002	Dec 17, 2016

VECURONIUM BROMIDE

INJECTABLE; INJECTION

VECURONIUM BROMIDE

<u>AP</u>	EUGIA PHARMA	<u>10MG/VIAL</u>	<u>A206670 001</u>	Dec 20, 2018
<u>AP</u>		<u>20MG/VIAL</u>	<u>A206670 002</u>	Dec 20, 2018
<u>AP</u>	GLAND	<u>10MG/VIAL</u>	<u>A205390 001</u>	May 26, 2016
<u>AP</u>		<u>20MG/VIAL</u>	<u>A205390 002</u>	May 26, 2016
<u>AP</u>	HIKMA	<u>10MG/VIAL</u>	<u>A075549 001</u>	Jun 13, 2000
<u>AP</u>		<u>10MG/VIAL</u>	<u>A203725 001</u>	Jul 30, 2019
<u>AP</u>		<u>20MG/VIAL</u>	<u>A075549 002</u>	Jun 13, 2000
<u>AP</u>		<u>20MG/VIAL</u>	<u>A203725 002</u>	Jul 30, 2019
<u>AP</u>	HOSPIRA	<u>10MG/VIAL</u>	<u>A075164 001</u>	Oct 21, 1999
<u>AP</u>		<u>20MG/VIAL</u>	<u>A075164 002</u>	Oct 21, 1999
<u>AP</u>	MEITHEAL	<u>10MG/VIAL</u>	<u>A074688 001</u>	Aug 25, 1999
<u>AP</u>		<u>20MG/VIAL</u>	<u>A074688 002</u>	Aug 25, 1999
<u>AP</u>	MYLAN LABS LTD	<u>10MG/VIAL</u>	<u>A090243 001</u>	May 11, 2010
<u>AP</u>		<u>20MG/VIAL</u>	<u>A090243 002</u>	May 11, 2010
<u>AP</u>	SAGENT PHARMS INC	<u>10MG/VIAL</u>	<u>A078274 001</u>	Dec 29, 2008
<u>AP</u>		<u>20MG/VIAL</u>	<u>A078274 002</u>	Dec 29, 2008
<u>AP</u>	! SUN PHARM	<u>10MG/VIAL</u>	<u>A079001 001</u>	Jun 17, 2009
<u>AP</u>	!	<u>20MG/VIAL</u>	<u>A079001 002</u>	Jun 17, 2009

VEMURAFENIB

TABLET; ORAL

ZELBORAF

+!	HOFFMANN LA ROCHE	240MG	N202429 001	Aug 17, 2011
----	-------------------	-------	-------------	--------------

VENETOCLAX

TABLET; ORAL

VENCLEXTA

+	ABBVIE	10MG	N208573 001	Apr 11, 2016
+		50MG	N208573 002	Apr 11, 2016
+!		100MG	N208573 003	Apr 11, 2016

VENLAFAXINE BESYLATE

TABLET, EXTENDED RELEASE; ORAL

VENLAFAXINE BESYLATE

+!	ALMATICA	EQ 112.5MG BASE	N215429 001	Jun 29, 2022
----	----------	-----------------	-------------	--------------

VENLAFAXINE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

EFFEXOR XR

<u>AB</u>	+	UPJOHN	<u>EQ 37.5MG BASE</u>	<u>N020699 001</u>	Oct 20, 1997
<u>AB</u>	+		<u>EQ 75MG BASE</u>	<u>N020699 002</u>	Oct 20, 1997
<u>AB</u>	+!		<u>EQ 150MG BASE</u>	<u>N020699 004</u>	Oct 20, 1997

VENLAFAXINE HYDROCHLORIDE

<u>AB</u>	ALEMBIC	<u>EQ 37.5MG BASE</u>	<u>A217767 001</u>	Jun 06, 2024
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A217767 002</u>	Jun 06, 2024
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A217767 003</u>	Jun 06, 2024
<u>AB</u>	ANNORA PHARMA	<u>EQ 37.5MG BASE</u>	<u>A212277 001</u>	Jul 08, 2019
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A212277 002</u>	Jul 08, 2019
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A212277 003</u>	Jul 08, 2019

PRESCRIPTION DRUG PRODUCT LIST

VENLAFAXINE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE;ORAL

VENLAFAXINE HYDROCHLORIDE

<u>AB</u>	AUROBINDO PHARMA LTD	<u>EQ 37.5MG BASE</u>	<u>A200834 001</u>	Apr 14, 2011
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A200834 002</u>	Apr 14, 2011
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A200834 003</u>	Apr 14, 2011
<u>AB</u>	DR REDDYS LABS LTD	<u>EQ 37.5MG BASE</u>	<u>A078421 001</u>	May 06, 2011
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A078421 002</u>	May 06, 2011
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A078421 003</u>	May 06, 2011
<u>AB</u>	GRANULES	<u>EQ 37.5MG BASE</u>	<u>A217390 001</u>	May 18, 2023
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A217390 002</u>	May 18, 2023
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A217390 003</u>	May 18, 2023
<u>AB</u>	INTELLIPHARMACEUTICS	<u>EQ 37.5MG BASE</u>	<u>A201272 001</u>	Nov 23, 2018
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A201272 002</u>	Nov 23, 2018
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A201272 003</u>	Nov 23, 2018
<u>AB</u>	INVENTIA HLTHCARE	<u>EQ 37.5MG BASE</u>	<u>A203332 001</u>	Mar 12, 2020
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A203332 002</u>	Mar 12, 2020
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A203332 003</u>	Mar 12, 2020
<u>AB</u>	MACLEODS PHARMS LTD	<u>EQ 37.5MG BASE</u>	<u>A204889 001</u>	Oct 05, 2017
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A204889 002</u>	Oct 05, 2017
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A204889 003</u>	Oct 05, 2017
<u>AB</u>	ORBION PHARMS	<u>EQ 37.5MG BASE</u>	<u>A091123 001</u>	Jul 11, 2011
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A091123 002</u>	Jul 11, 2011
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A091123 003</u>	Jul 11, 2011
<u>AB</u>	TEVA	<u>EQ 37.5MG BASE</u>	<u>A076565 001</u>	Jun 28, 2010
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A076565 002</u>	Jun 28, 2010
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A076565 003</u>	Jun 28, 2010
<u>AB</u>	WOCKHARDT BIO AG	<u>EQ 37.5MG BASE</u>	<u>A078865 001</u>	Apr 14, 2011
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A078865 002</u>	Apr 14, 2011
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A078865 003</u>	Apr 14, 2011
<u>AB</u>	YICHANG HUMANWELL	<u>EQ 37.5MG BASE</u>	<u>A214654 001</u>	Aug 06, 2021
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A214654 002</u>	Aug 06, 2021
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A214654 003</u>	Aug 06, 2021
<u>AB</u>	ZYDUS PHARMS USA INC	<u>EQ 37.5MG BASE</u>	<u>A090174 001</u>	Apr 14, 2011
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A090174 002</u>	Apr 14, 2011
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A090174 003</u>	Apr 14, 2011

TABLET;ORAL

VENLAFAXINE HYDROCHLORIDE

<u>AB</u>	ALEMbic PHARMS LTD	<u>EQ 25MG BASE</u>	<u>A078932 001</u>	Dec 14, 2010
<u>AB</u>		<u>EQ 37.5MG BASE</u>	<u>A078932 002</u>	Dec 14, 2010
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A078932 003</u>	Dec 14, 2010
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A078932 004</u>	Dec 14, 2010
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A078932 005</u>	Dec 14, 2010
<u>AB</u>	AUROBINDO PHARMA	<u>EQ 25MG BASE</u>	<u>A090555 001</u>	Apr 07, 2010
<u>AB</u>		<u>EQ 37.5MG BASE</u>	<u>A090555 002</u>	Apr 07, 2010
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A090555 003</u>	Apr 07, 2010
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A090555 004</u>	Apr 07, 2010
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A090555 005</u>	Apr 07, 2010
<u>AB</u>	CADILA PHARMS LTD	<u>EQ 25MG BASE</u>	<u>A206250 001</u>	Nov 21, 2018
<u>AB</u>		<u>EQ 37.5MG BASE</u>	<u>A206250 002</u>	Nov 21, 2018
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A206250 003</u>	Nov 21, 2018
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A206250 004</u>	Nov 21, 2018
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A206250 005</u>	Nov 21, 2018
<u>AB</u>	DR REDDYS LABS LTD	<u>EQ 25MG BASE</u>	<u>A078301 001</u>	Jun 13, 2008
<u>AB</u>		<u>EQ 37.5MG BASE</u>	<u>A078301 002</u>	Jun 13, 2008
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A078301 003</u>	Jun 13, 2008
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A078301 004</u>	Jun 13, 2008
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A078301 005</u>	Jun 13, 2008
<u>AB</u>	HERITAGE	<u>EQ 25MG BASE</u>	<u>A078554 001</u>	Jan 09, 2009
<u>AB</u>		<u>EQ 37.5MG BASE</u>	<u>A078554 002</u>	Jan 09, 2009
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A078554 003</u>	Jan 09, 2009
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A078554 004</u>	Jan 09, 2009
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A078554 005</u>	Jan 09, 2009
<u>AB</u>	SUN PHARM INDS INC	<u>EQ 25MG BASE</u>	<u>A078627 001</u>	Jun 13, 2008
<u>AB</u>		<u>EQ 37.5MG BASE</u>	<u>A078627 002</u>	Jun 13, 2008
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A078627 003</u>	Jun 13, 2008
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A078627 004</u>	Jun 13, 2008
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A078627 005</u>	Jun 13, 2008
<u>AB</u>	TEVA	<u>EQ 25MG BASE</u>	<u>A076690 001</u>	Aug 03, 2006
<u>AB</u>		<u>EQ 37.5MG BASE</u>	<u>A076690 002</u>	Aug 03, 2006

PRESCRIPTION DRUG PRODUCT LIST

VENLAFAXINE HYDROCHLORIDE

TABLET; ORAL

VENLAFAXINE HYDROCHLORIDE

<u>AB</u>	!	<u>EQ 50MG BASE</u>	<u>A076690 003</u>	Aug 03, 2006
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A076690 004</u>	Aug 03, 2006
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A076690 005</u>	Aug 03, 2006
<u>AB</u>	YAOPHARMA CO LTD	<u>EQ 25MG BASE</u>	<u>A202036 001</u>	May 28, 2015
<u>AB</u>		<u>EQ 37.5MG BASE</u>	<u>A202036 002</u>	May 28, 2015
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A202036 003</u>	May 28, 2015
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A202036 004</u>	May 28, 2015
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A202036 005</u>	May 28, 2015
<u>AB</u>	ZYDUS PHARMS USA	<u>EQ 25MG BASE</u>	<u>A077653 001</u>	Jun 13, 2008
<u>AB</u>		<u>EQ 37.5MG BASE</u>	<u>A077653 002</u>	Jun 13, 2008
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A077653 003</u>	Jun 13, 2008
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A077653 004</u>	Jun 13, 2008
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A077653 005</u>	Jun 13, 2008

TABLET, EXTENDED RELEASE; ORAL

VENLAFAXINE HYDROCHLORIDE

<u>AB</u>	AJANTA PHARMA LTD	<u>EQ 37.5MG BASE</u>	<u>A214691 001</u>	Apr 12, 2023	
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A214691 002</u>	Apr 12, 2023	
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A214691 003</u>	Apr 12, 2023	
<u>AB</u>		<u>EQ 225MG BASE</u>	<u>A214691 004</u>	Apr 12, 2023	
<u>AB</u>	ALKEM LABS LTD	<u>EQ 37.5MG BASE</u>	<u>A214127 001</u>	Jun 10, 2021	
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A214127 002</u>	Jun 10, 2021	
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A214127 003</u>	Jun 10, 2021	
<u>AB</u>		<u>EQ 225MG BASE</u>	<u>A214127 004</u>	Jun 10, 2021	
<u>AB</u>	APPCO	<u>EQ 150MG BASE</u>	<u>A214609 001</u>	Jun 30, 2021	
<u>AB</u>		<u>EQ 225MG BASE</u>	<u>A214609 002</u>	Jun 30, 2021	
<u>AB</u>	ASCENT PHARMS INC	<u>EQ 37.5MG BASE</u>	<u>A214419 001</u>	Oct 21, 2020	
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A214419 002</u>	Oct 21, 2020	
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A214419 003</u>	Oct 21, 2020	
<u>AB</u>		<u>EQ 225MG BASE</u>	<u>A214419 004</u>	Oct 21, 2020	
<u>AB</u>	CADILA PHARMS LTD	<u>EQ 75MG BASE</u>	<u>A211323 001</u>	Aug 29, 2019	
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A211323 002</u>	Aug 29, 2019	
<u>AB</u>		<u>EQ 225MG BASE</u>	<u>A211323 003</u>	Aug 29, 2019	
<u>AB</u>	DEXCEL	<u>EQ 75MG BASE</u>	<u>A213927 001</u>	Jan 21, 2021	
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A209193 001</u>	Oct 31, 2019	
<u>AB</u>		<u>EQ 225MG BASE</u>	<u>A209193 002</u>	Oct 31, 2019	
<u>AB</u>	NOSTRUM LABS INC	<u>EQ 150MG BASE</u>	<u>A205468 002</u>	Mar 24, 2017	
<u>AB</u>		<u>EQ 225MG BASE</u>	<u>A205468 003</u>	Mar 24, 2017	
<u>AB</u>	+	<u>OSMOTICA PHARM US</u>	<u>EQ 37.5MG BASE</u>	<u>N022104 001</u>	May 20, 2008
<u>AB</u>	+	<u>EQ 75MG BASE</u>	<u>N022104 002</u>	May 20, 2008	
<u>AB</u>	+	<u>EQ 150MG BASE</u>	<u>N022104 003</u>	May 20, 2008	
<u>AB</u>	+	<u>EQ 225MG BASE</u>	<u>N022104 004</u>	May 20, 2008	
<u>AB</u>	UNIQUE	<u>EQ 37.5MG BASE</u>	<u>A216044 001</u>	Nov 28, 2022	
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A216044 002</u>	Nov 28, 2022	
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A216044 003</u>	Nov 28, 2022	
<u>AB</u>		<u>EQ 225MG BASE</u>	<u>A216044 004</u>	Nov 28, 2022	
<u>AB</u>	ZYDUS PHARMS	<u>EQ 37.5MG BASE</u>	<u>A215622 001</u>	Aug 30, 2022	
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A215622 002</u>	Aug 30, 2022	
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A215622 003</u>	Aug 30, 2022	
<u>AB</u>		<u>EQ 225MG BASE</u>	<u>A215622 004</u>	Aug 30, 2022	

VERAPAMIL HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

VERAPAMIL HYDROCHLORIDE

<u>AB</u>	MYLAN	<u>120MG</u>	<u>A075138 001</u>	Apr 20, 1999	
<u>AB</u>		<u>180MG</u>	<u>A075138 002</u>	Apr 20, 1999	
<u>AB</u>		<u>240MG</u>	<u>A075138 003</u>	Apr 20, 1999	
<u>VERELAN</u>					
<u>AB</u>	+	<u>AZURITY</u>	<u>120MG</u>	<u>N019614 001</u>	May 29, 1990
<u>AB</u>	+	<u>180MG</u>	<u>N019614 003</u>	Jan 09, 1992	
<u>AB</u>	+	<u>240MG</u>	<u>N019614 002</u>	May 29, 1990	
	+	<u>360MG</u>	<u>N019614 004</u>	May 10, 1996	
VERELAN PM					
	+	<u>AZURITY</u>	<u>100MG</u>	<u>N020943 001</u>	Nov 25, 1998
	+	<u>200MG</u>	<u>N020943 002</u>	Nov 25, 1998	
	+	<u>300MG</u>	<u>N020943 003</u>	Nov 25, 1998	

INJECTABLE; INJECTION

VERAPAMIL HYDROCHLORIDE

<u>AP</u>	!	EUGIA PHARMA	<u>2.5MG/ML</u>	<u>A212965 001</u>	Jul 06, 2020
<u>AP</u>		FRESENIUS KABI USA	<u>2.5MG/ML</u>	<u>A216471 001</u>	Apr 23, 2024
<u>AP</u>		GLAND PHARMA LTD	<u>2.5MG/ML</u>	<u>A214361 001</u>	Oct 15, 2020

PRESCRIPTION DRUG PRODUCT LIST

VERAPAMIL HYDROCHLORIDE

INJECTABLE; INJECTION

VERAPAMIL HYDROCHLORIDE

<u>AP</u>	HERITAGE	<u>2.5MG/ML</u>	<u>A215829</u>	<u>001</u>	Mar 17, 2022
<u>AP</u>	MANKIND PHARMA	<u>2.5MG/ML</u>	<u>A214653</u>	<u>001</u>	Jun 15, 2022
<u>AP</u>	ZYDUS PHARMS	<u>2.5MG/ML</u>	<u>A214215</u>	<u>001</u>	Oct 15, 2020

SOLUTION; INTRAVENOUS

VERAPAMIL HYDROCHLORIDE

<u>AP</u>	AMNEAL	<u>5MG/2ML (2.5MG/ML)</u>	<u>A210994</u>	<u>001</u>	Jul 13, 2018
<u>AP</u>		<u>10MG/4ML (2.5MG/ML)</u>	<u>A210994</u>	<u>002</u>	Jul 13, 2018
<u>AP</u>	CAPLIN	<u>5MG/2ML (2.5MG/ML)</u>	<u>A213232</u>	<u>001</u>	Mar 25, 2020
<u>AP</u>		<u>10MG/4ML (2.5MG/ML)</u>	<u>A213232</u>	<u>002</u>	Mar 25, 2020
<u>AP</u>	EXELA PHARMA	<u>5MG/2ML (2.5MG/ML)</u>	<u>N018925</u>	<u>001</u>	Mar 30, 1984
<u>AP</u>	! HOSPIRA	<u>5MG/2ML (2.5MG/ML)</u>	<u>A070738</u>	<u>001</u>	May 06, 1987
<u>AP</u>	!	<u>5MG/2ML (2.5MG/ML)</u>	<u>A075136</u>	<u>001</u>	Oct 20, 1998
<u>AP</u>	!	<u>5MG/2ML (2.5MG/ML)</u>	<u>A070737</u>	<u>001</u>	May 06, 1987
<u>AP</u>	!	<u>10MG/4ML (2.5MG/ML)</u>	<u>A070737</u>	<u>002</u>	May 06, 1987
<u>AP</u>	MICRO LABS	<u>5MG/2ML (2.5MG/ML)</u>	<u>A211370</u>	<u>001</u>	Dec 28, 2018
<u>AP</u>		<u>10MG/4ML (2.5MG/ML)</u>	<u>A211370</u>	<u>002</u>	Dec 28, 2018
<u>AP</u>	NEPHRON	<u>5MG/2ML (2.5MG/ML)</u>	<u>A213352</u>	<u>002</u>	Sep 16, 2020
<u>AP</u>		<u>10MG/4ML (2.5MG/ML)</u>	<u>A213352</u>	<u>001</u>	Mar 17, 2020
<u>AP</u>	SOMERSET	<u>5MG/2ML (2.5MG/ML)</u>	<u>A211035</u>	<u>001</u>	Jun 18, 2018
<u>AP</u>		<u>10MG/4ML (2.5MG/ML)</u>	<u>A211035</u>	<u>002</u>	Jun 18, 2018
<u>AP</u>	SOMERSET THERAPS LLC	<u>5MG/2ML (2.5MG/ML)</u>	<u>A211015</u>	<u>001</u>	Jun 18, 2018
<u>AP</u>		<u>10MG/4ML (2.5MG/ML)</u>	<u>A211015</u>	<u>002</u>	Jun 18, 2018

TABLET; ORAL

VERAPAMIL HYDROCHLORIDE

<u>AB</u>	HERITAGE	<u>40MG</u>	<u>A071881</u>	<u>002</u>	Oct 14, 2015
<u>AB</u>		<u>80MG</u>	<u>A071881</u>	<u>003</u>	Apr 05, 1988
<u>AB</u>	!	<u>120MG</u>	<u>A071881</u>	<u>001</u>	Apr 05, 1988
<u>AB</u>	WATSON LABS	<u>40MG</u>	<u>A072924</u>	<u>001</u>	Jun 29, 1993
<u>AB</u>		<u>80MG</u>	<u>A070995</u>	<u>001</u>	Oct 01, 1986
<u>AB</u>		<u>120MG</u>	<u>A070994</u>	<u>001</u>	Oct 01, 1986

TABLET, EXTENDED RELEASE; ORAL

VERAPAMIL HYDROCHLORIDE

<u>AB</u>	CADILA PHARMS LTD	<u>120MG</u>	<u>A206173</u>	<u>003</u>	Nov 14, 2022
<u>AB</u>		<u>180MG</u>	<u>A206173</u>	<u>001</u>	May 05, 2017
<u>AB</u>		<u>240MG</u>	<u>A206173</u>	<u>002</u>	May 05, 2017
<u>AB</u>	! GLENMARK PHARMS LTD	<u>120MG</u>	<u>A090700</u>	<u>001</u>	Aug 03, 2011
<u>AB</u>	!	<u>180MG</u>	<u>A090700</u>	<u>002</u>	Aug 03, 2011
<u>AB</u>	!	<u>240MG</u>	<u>A078906</u>	<u>001</u>	Sep 17, 2009
<u>AB</u>	STRIDES PHARMA	<u>120MG</u>	<u>A075072</u>	<u>001</u>	May 25, 1999
<u>AB</u>		<u>240MG</u>	<u>A075072</u>	<u>003</u>	May 25, 1999

VERICIGUAT

TABLET; ORAL

VERQUVO

+	MSD	2.5MG	N214377	001	Jan 19, 2021
+		5MG	N214377	002	Jan 19, 2021
+	!	10MG	N214377	003	Jan 19, 2021

VERTEPORFIN

INJECTABLE; INJECTION

VISUDYNE

+	!	BAUSCH LOMB IRELAND	15MG/VIAL	N021119	001	Apr 12, 2000
---	---	---------------------	-----------	---------	-----	--------------

VIBEGRON

TABLET; ORAL

GEMTESA

+	!	UROVANT	75MG	N213006	001	Dec 23, 2020
---	---	---------	------	---------	-----	--------------

VIGABATRIN

FOR SOLUTION; ORAL

SABRIL

<u>AA</u>	+	!	LUNDBECK PHARMS LLC	<u>500MG/PACKET</u>	<u>N022006</u>	<u>001</u>	Aug 21, 2009
-----------	---	---	---------------------	---------------------	----------------	------------	--------------

VIGABATRIN

<u>AA</u>		ALKEM LABS LTD	<u>500MG/PACKET</u>	<u>A213375</u>	<u>001</u>	Dec 02, 2020
<u>AA</u>		AMNEAL PHARMS	<u>500MG/PACKET</u>	<u>A210155</u>	<u>001</u>	Mar 13, 2018
<u>AA</u>		ANNORA PHARMA	<u>500MG/PACKET</u>	<u>A213519</u>	<u>001</u>	Jan 26, 2021
<u>AA</u>		AUROBINDO PHARMA LTD	<u>500MG/PACKET</u>	<u>A213899</u>	<u>001</u>	Sep 29, 2021
<u>AA</u>		DEXCEL	<u>500MG/PACKET</u>	<u>A214992</u>	<u>001</u>	May 13, 2021
<u>AA</u>		DR REDDYS	<u>500MG/PACKET</u>	<u>A211481</u>	<u>001</u>	Nov 20, 2018
<u>AA</u>		ENDO OPERATIONS	<u>500MG/PACKET</u>	<u>A208218</u>	<u>001</u>	Apr 27, 2017

PRESCRIPTION DRUG PRODUCT LIST

VIGABATRIN

FOR SOLUTION;ORAL

VIGABATRIN

AA	INVAGEN PHARMS	500MG/PACKET	A211592 001	Dec 03, 2019
AA	MSN	500MG/PACKET	A215363 001	Sep 07, 2022
AA	TEVA PHARMS USA	500MG/PACKET	A209824 001	Apr 23, 2018

VIGADRONE

AA	AUCTA	500MG/PACKET	A210196 001	Jun 21, 2018
-----------	-------	---------------------	--------------------	--------------

VIGPODER

AA	PYROS PHARMS	500MG/PACKET	A214961 001	Jun 24, 2022
-----------	--------------	---------------------	--------------------	--------------

SOLUTION;ORAL

VIGAFYDE

+	!	PYROS PHARMS	100MG/ML	N217684 001	Jun 17, 2024
---	---	--------------	----------	-------------	--------------

TABLET;ORAL

SABRII

AB	+	!	LUNDBECK PHARMS LLC	500MG	N020427 001	Aug 21, 2009
-----------	---	---	---------------------	--------------	--------------------	--------------

VIGABATRIN

AB	AMNEAL PHARMS	500MG	A210042 001	Jun 22, 2022
AB	AUROBINDO PHARMA LTD	500MG	A215601 001	May 10, 2022
AB	DEXCEL	500MG	A215109 001	Sep 23, 2021
AB	DR REDDYS	500MG	A211539 001	Jan 29, 2021
AB	HIKMA	500MG	A213104 001	Aug 29, 2022
AB	RYAN LABS	500MG	A215519 001	Apr 28, 2023
AB	TEVA PHARMS USA	500MG	A209822 001	Jan 14, 2019
AB	UPSHER SMITH LABS	500MG	A214749 001	Jun 29, 2023
AB	ZYDUS LIFESCIENCES	500MG	A215707 001	Jan 19, 2022

VILAZODONE HYDROCHLORIDE

TABLET;ORAL

VIIBRYD

AB	+	!	ABEVIE	10MG	N022567 001	Jan 21, 2011
AB	+			20MG	N022567 002	Jan 21, 2011
AB	+			40MG	N022567 003	Jan 21, 2011

VILAZODONE HYDROCHLORIDE

AB	ACCORD HLTHCARE	10MG	A208209 001	Apr 27, 2021
AB		20MG	A208209 002	Apr 27, 2021
AB		40MG	A208209 003	Apr 27, 2021
AB	ALEMBIC	10MG	A208202 001	Jan 10, 2020
AB		20MG	A208202 002	Jan 10, 2020
AB		40MG	A208202 003	Jan 10, 2020
AB	APOTEX	10MG	A208228 001	Jul 07, 2023
AB		20MG	A208228 002	Jul 07, 2023
AB		40MG	A208228 003	Jul 07, 2023
AB	INVAGEN PHARMS	10MG	A208200 001	Apr 07, 2021
AB		20MG	A208200 003	Dec 06, 2022
AB		40MG	A208200 002	Apr 07, 2021
AB	TEVA PHARMS USA	10MG	A208212 001	Sep 30, 2019
AB		20MG	A208212 002	Sep 30, 2019
AB		40MG	A208212 003	Sep 30, 2019

VILOXAZINE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE;ORAL

QELBREE

+		SUPERUS PHARMS	EQ 100MG BASE	N211964 001	Apr 02, 2021
+			EQ 150MG BASE	N211964 002	Apr 02, 2021
+	!		EQ 200MG BASE	N211964 003	Apr 02, 2021

VILTOLARSEN

SOLUTION;INTRAVENOUS

VILTEPSO

+	!	NIPPON SHINYAKU	250MG/5ML (50MG/ML)	N212154 001	Aug 12, 2020
---	---	-----------------	---------------------	-------------	--------------

VINBLASTINE SULFATE

INJECTABLE;INJECTION

VINBLASTINE SULFATE

!		FRESENIUS KABI USA	1MG/ML	A089515 001	Apr 29, 1987
!		HIKMA	10MG/VIAL	A089395 001	Apr 09, 1987

PRESCRIPTION DRUG PRODUCT LIST

VINCRISTINE SULFATE

INJECTABLE; INJECTION

VINCRISTINE SULFATE PFS

! HOSPIRA

1MG/ML

A071484 001 Apr 19, 1988

VINORELBINE TARTRATE

INJECTABLE; INJECTION

VINORELBINE TARTRATEAP ACTAVIS TOTOWAEQ 10MG BASE/MLA078011 001 Jul 22, 2009AP DR REDDYSEQ 10MG BASE/MLA202017 001 Sep 12, 2013AP HIKMAEQ 10MG BASE/MLA075992 001 Jun 10, 2003APEQ 10MG BASE/MLA076461 001 Dec 11, 2003AP ! JIANGSU HANSOH
PHARMEQ 10MG BASE/MLA091106 001 Sep 26, 2012VISMODEGIB

CAPSULE; ORAL

ERIVEDGE

+! GENENTECH

150MG

N203388 001 Jan 30, 2012

VITAMIN A PALMITATE

INJECTABLE; INJECTION

AQUASOL A

+! CASPER PHARMA LLC

EQ 50,000 UNITS BASE/ML

N006823 001

VOCLOSPORIN

CAPSULE; ORAL

LUPKYNIS

+! AURINIA

7.9MG

N213716 001 Jan 22, 2021

VONOPRAZAN FUMARATE

TABLET; ORAL

VOQUEZNA

+ PHATHOM

EQ 10MG BASE

N215151 001 Nov 01, 2023

+!

EQ 20MG BASE

N215151 002 Nov 01, 2023

VORASIDENIB

TABLET; ORAL

VORANIGO

+ SERVIER

10MG

N218784 001 Aug 06, 2024

+!

40MG

N218784 002 Aug 06, 2024

VORICONAZOLE

FOR SUSPENSION; ORAL

VFENDAB +! PF PRISM CV200MG/5MLN021630 001 Dec 19, 2003VORICONAZOLEAB AMNEAL PHARMS200MG/5MLA205034 001 Apr 13, 2016AB HAINAN POLY200MG/5MLA216805 001 Jan 02, 2024AB NOVEL LABS INC200MG/5MLA206799 001 May 31, 2016

INJECTABLE; INTRAVENOUS

VFENDAP +! PF PRISM CV200MG/VIALN021267 001 May 24, 2002VORICONAZOLEAP ALMAJECT200MG/VIALA206398 001 Mar 23, 2016AP ASPIRO200MG/VIALA218533 001 Apr 03, 2024AP GLAND PHARMA LTD200MG/VIALA211099 001 Mar 31, 2020AP HIKMA200MG/VIALN208562 001 Mar 09, 2017AP MEITHEAL200MG/VIALA214516 001 May 09, 2022AP SANDOZ INC200MG/VIALA090862 001 May 30, 2012AP SLATE RUN PHARMA200MG/VIALA211661 001 Nov 30, 2018AP UBI200MG/VIALA211264 001 Mar 09, 2023AP ZYDUS PHARMS200MG/VIALA208983 001 Jul 16, 2018

TABLET; ORAL

VFENDAB + PF PRISM CV50MGN021266 001 May 24, 2002AB +!200MGN021266 002 May 24, 2002VORICONAZOLEAB AJANTA PHARMA LTD50MGA206181 001 May 24, 2016AB200MGA206181 002 May 24, 2016AB AUROBINDO PHARMA50MGA206837 001 Jan 22, 2016

LTD

AB200MGA206837 002 Jan 22, 2016AB CADILA50MGA206747 001 May 24, 2016AB200MGA206747 002 May 24, 2016AB CHARTWELL RX50MGA207371 001 May 24, 2016

PRESCRIPTION DRUG PRODUCT LIST

VORICONAZOLE

TABLET; ORAL

VORICONAZOLE

<u>AB</u>		<u>200MG</u>	<u>A207371</u>	<u>002</u>	May 24, 2016
<u>AB</u>	EPIC PHARMA LLC	<u>50MG</u>	<u>A207049</u>	<u>001</u>	Sep 07, 2016
<u>AB</u>		<u>200MG</u>	<u>A207049</u>	<u>002</u>	Sep 07, 2016
<u>AB</u>	GLENMARK PHARMS LTD	<u>50MG</u>	<u>A203503</u>	<u>001</u>	Sep 02, 2015
<u>AB</u>		<u>200MG</u>	<u>A203503</u>	<u>002</u>	Sep 02, 2015
<u>AB</u>	MYLAN PHARMS INC	<u>50MG</u>	<u>A090547</u>	<u>001</u>	Apr 22, 2010
<u>AB</u>		<u>200MG</u>	<u>A090547</u>	<u>002</u>	Apr 22, 2010
<u>AB</u>	PRINSTON INC	<u>50MG</u>	<u>A206654</u>	<u>001</u>	Aug 08, 2016
<u>AB</u>		<u>200MG</u>	<u>A206654</u>	<u>002</u>	Aug 08, 2016
<u>AB</u>	RISING	<u>50MG</u>	<u>A206762</u>	<u>001</u>	May 24, 2016
<u>AB</u>		<u>200MG</u>	<u>A206762</u>	<u>002</u>	May 24, 2016
<u>AB</u>	SANDOZ INC	<u>50MG</u>	<u>A200265</u>	<u>001</u>	Dec 12, 2011
<u>AB</u>		<u>200MG</u>	<u>A200265</u>	<u>002</u>	Dec 12, 2011

VORINOSTAT

CAPSULE; ORAL

ZOLINZA

+! MSD SUB MERCK 100MG N021991 001 Oct 06, 2006

VORTIOXETINE HYDROBROMIDE

TABLET; ORAL

TRINTELLIX

+	TAKEDA PHARMS USA	EQ 5MG BASE	N204447	001	Sep 30, 2013
+		EQ 10MG BASE	N204447	002	Sep 30, 2013
+!		EQ 20MG BASE	N204447	004	Sep 30, 2013

VOSORITIDE

POWDER; SUBCUTANEOUS

VOXZOGO

+	BIOMARIN PHARM	0.4MG/VIAL	N214938	001	Nov 19, 2021
+		0.56MG/VIAL	N214938	002	Nov 19, 2021
+		1.2MG/VIAL	N214938	003	Nov 19, 2021

VUTRISIRAN SODIUM

SOLUTION; SUBCUTANEOUS

AMVUTTRA

+! ALNYLAM PHARMS INC EQ 25MG BASE/0.5ML (EQ 25MG BASE/0.5ML) N215515 001 Jun 13, 2022

WARFARIN SODIUM

TABLET; ORAL

JANTOVEN

<u>AB</u>	UPSHER SMITH LABS	<u>1MG</u>	<u>A040416</u>	<u>001</u>	Oct 02, 2003
<u>AB</u>		<u>2MG</u>	<u>A040416</u>	<u>002</u>	Oct 02, 2003
<u>AB</u>		<u>2.5MG</u>	<u>A040416</u>	<u>003</u>	Oct 02, 2003
<u>AB</u>		<u>3MG</u>	<u>A040416</u>	<u>004</u>	Oct 02, 2003
<u>AB</u>		<u>4MG</u>	<u>A040416</u>	<u>005</u>	Oct 02, 2003
<u>AB</u>		<u>5MG</u>	<u>A040416</u>	<u>006</u>	Oct 02, 2003
<u>AB</u>		<u>6MG</u>	<u>A040416</u>	<u>007</u>	Oct 02, 2003
<u>AB</u>		<u>7.5MG</u>	<u>A040416</u>	<u>008</u>	Oct 02, 2003
<u>AB</u>		<u>10MG</u>	<u>A040416</u>	<u>009</u>	Oct 02, 2003

WARFARIN SODIUM

<u>AB</u>	AMNEAL PHARMS	<u>1MG</u>	<u>A202202</u>	<u>001</u>	Mar 04, 2013
<u>AB</u>		<u>2MG</u>	<u>A202202</u>	<u>002</u>	Mar 04, 2013
<u>AB</u>		<u>2.5MG</u>	<u>A202202</u>	<u>003</u>	Mar 04, 2013
<u>AB</u>		<u>3MG</u>	<u>A202202</u>	<u>004</u>	Mar 04, 2013
<u>AB</u>		<u>4MG</u>	<u>A202202</u>	<u>005</u>	Mar 04, 2013
<u>AB</u>		<u>5MG</u>	<u>A202202</u>	<u>006</u>	Mar 04, 2013
<u>AB</u>		<u>6MG</u>	<u>A202202</u>	<u>007</u>	Mar 04, 2013
<u>AB</u>		<u>7.5MG</u>	<u>A202202</u>	<u>008</u>	Mar 04, 2013
<u>AB</u>		<u>10MG</u>	<u>A202202</u>	<u>009</u>	Mar 04, 2013
<u>AB</u>	INVAGEN PHARMS	<u>1MG</u>	<u>A090935</u>	<u>001</u>	May 25, 2011
<u>AB</u>		<u>2MG</u>	<u>A090935</u>	<u>002</u>	May 25, 2011
<u>AB</u>		<u>2.5MG</u>	<u>A090935</u>	<u>003</u>	May 25, 2011
<u>AB</u>		<u>3MG</u>	<u>A090935</u>	<u>004</u>	May 25, 2011
<u>AB</u>		<u>4MG</u>	<u>A090935</u>	<u>005</u>	May 25, 2011
<u>AB</u>		<u>5MG</u>	<u>A090935</u>	<u>006</u>	May 25, 2011
<u>AB</u>		<u>6MG</u>	<u>A090935</u>	<u>007</u>	May 25, 2011
<u>AB</u>		<u>7.5MG</u>	<u>A090935</u>	<u>008</u>	May 25, 2011
<u>AB</u>		<u>10MG</u>	<u>A090935</u>	<u>009</u>	May 25, 2011
<u>AB</u>	PLIVA	<u>1MG</u>	<u>A040616</u>	<u>009</u>	Jul 05, 2006
<u>AB</u>		<u>2MG</u>	<u>A040616</u>	<u>001</u>	Jul 05, 2006
<u>AB</u>		<u>2.5MG</u>	<u>A040616</u>	<u>002</u>	Jul 05, 2006

PRESCRIPTION DRUG PRODUCT LIST

WARFARIN SODIUM

TABLET; ORAL

WARFARIN SODIUM

<u>AB</u>		<u>3MG</u>	<u>A040616</u>	<u>003</u>	Jul 05, 2006
<u>AB</u>		<u>4MG</u>	<u>A040616</u>	<u>004</u>	Jul 05, 2006
<u>AB</u>		<u>5MG</u>	<u>A040616</u>	<u>005</u>	Jul 05, 2006
<u>AB</u>		<u>6MG</u>	<u>A040616</u>	<u>006</u>	Jul 05, 2006
<u>AB</u>		<u>7.5MG</u>	<u>A040616</u>	<u>007</u>	Jul 05, 2006
<u>AB</u>	!	<u>10MG</u>	<u>A040616</u>	<u>008</u>	Jul 05, 2006
<u>AB</u>	TARO	<u>1MG</u>	<u>A040301</u>	<u>002</u>	Jul 15, 1999
<u>AB</u>		<u>2MG</u>	<u>A040301</u>	<u>003</u>	Jul 15, 1999
<u>AB</u>		<u>2.5MG</u>	<u>A040301</u>	<u>004</u>	Jul 15, 1999
<u>AB</u>		<u>3MG</u>	<u>A040301</u>	<u>005</u>	Jul 15, 1999
<u>AB</u>		<u>4MG</u>	<u>A040301</u>	<u>006</u>	Jul 15, 1999
<u>AB</u>		<u>5MG</u>	<u>A040301</u>	<u>007</u>	Jul 15, 1999
<u>AB</u>		<u>6MG</u>	<u>A040301</u>	<u>008</u>	Jul 15, 1999
<u>AB</u>		<u>7.5MG</u>	<u>A040301</u>	<u>009</u>	Jul 15, 1999
<u>AB</u>		<u>10MG</u>	<u>A040301</u>	<u>001</u>	Jul 15, 1999
<u>AB</u>	ZYDUS PHARMS USA	<u>1MG</u>	<u>A040663</u>	<u>001</u>	May 30, 2006
<u>AB</u>		<u>2MG</u>	<u>A040663</u>	<u>002</u>	May 30, 2006
<u>AB</u>		<u>2.5MG</u>	<u>A040663</u>	<u>003</u>	May 30, 2006
<u>AB</u>		<u>3MG</u>	<u>A040663</u>	<u>004</u>	May 30, 2006
<u>AB</u>		<u>4MG</u>	<u>A040663</u>	<u>005</u>	May 30, 2006
<u>AB</u>		<u>5MG</u>	<u>A040663</u>	<u>006</u>	May 30, 2006
<u>AB</u>		<u>6MG</u>	<u>A040663</u>	<u>007</u>	May 30, 2006
<u>AB</u>		<u>7.5MG</u>	<u>A040663</u>	<u>008</u>	May 30, 2006
<u>AB</u>		<u>10MG</u>	<u>A040663</u>	<u>009</u>	May 30, 2006

XENON XE-129 HYPERPOLARIZED

GAS; INHALATION

XENOVIEW

+! POLAREAN

N/A

N214375 001 Dec 23, 2022

XENON XE-133

GAS; INHALATION

XENON XE 133

CURIUM

10mCi/VIAL

N018327 001 Mar 09, 1982

20mCi/VIAL

N018327 002 Mar 09, 1982

LANTHEUS MEDCL

10mCi/VIAL

N017284 001

20mCi/VIAL

N017284 002

ZAFIRLUKAST

TABLET; ORAL

ACCOLATE

<u>AB</u>	+	STRIDES PHARMA	<u>10MG</u>	<u>N020547</u>	<u>003</u>	Sep 17, 1999
<u>AB</u>	+	!	<u>20MG</u>	<u>N020547</u>	<u>001</u>	Sep 26, 1996

ZAFIRLUKAST

<u>AB</u>		ANNORA PHARMA	<u>10MG</u>	<u>A212475</u>	<u>001</u>	Sep 10, 2020
<u>AB</u>			<u>20MG</u>	<u>A212475</u>	<u>002</u>	Sep 10, 2020
<u>AB</u>		AUROBINDO PHARMA	<u>10MG</u>	<u>A213163</u>	<u>001</u>	Nov 27, 2023
<u>AB</u>			<u>20MG</u>	<u>A213163</u>	<u>002</u>	Nov 27, 2023
<u>AB</u>		DR REDDYS LABS LTD	<u>10MG</u>	<u>A090372</u>	<u>001</u>	Nov 18, 2010
<u>AB</u>			<u>20MG</u>	<u>A090372</u>	<u>002</u>	Nov 18, 2010
<u>AB</u>		RISING PHARMS	<u>10MG</u>	<u>A204928</u>	<u>001</u>	Aug 25, 2022
<u>AB</u>			<u>20MG</u>	<u>A204928</u>	<u>002</u>	Aug 25, 2022

ZALEPLON

CAPSULE; ORAL

SONATA

<u>AB</u>	+	PFIZER	<u>5MG</u>	<u>N020859</u>	<u>001</u>	Aug 13, 1999
<u>AB</u>	+	!	<u>10MG</u>	<u>N020859</u>	<u>002</u>	Aug 13, 1999

ZALEPLON

<u>AB</u>		AUROBINDO PHARMA	<u>5MG</u>	<u>A078829</u>	<u>001</u>	Jun 06, 2008
<u>AB</u>			<u>10MG</u>	<u>A078829</u>	<u>002</u>	Jun 06, 2008
<u>AB</u>		CHARTWELL MOLECULAR	<u>5MG</u>	<u>A077505</u>	<u>001</u>	Jun 20, 2008
<u>AB</u>			<u>10MG</u>	<u>A077505</u>	<u>002</u>	Jun 20, 2008
<u>AB</u>		HIKMA	<u>5MG</u>	<u>A077237</u>	<u>001</u>	Jun 06, 2008
<u>AB</u>			<u>10MG</u>	<u>A077237</u>	<u>002</u>	Jun 06, 2008
<u>AB</u>		ORBION PHARMS	<u>5MG</u>	<u>A090374</u>	<u>001</u>	Sep 17, 2009
<u>AB</u>			<u>10MG</u>	<u>A090374</u>	<u>002</u>	Sep 17, 2009
<u>AB</u>		UNICHEM	<u>5MG</u>	<u>A078989</u>	<u>001</u>	Jun 06, 2008
<u>AB</u>			<u>10MG</u>	<u>A078989</u>	<u>002</u>	Jun 06, 2008

PRESCRIPTION DRUG PRODUCT LIST

ZANAMIVIR

POWDER; INHALATION

RELENZA

+! GLAXOSMITHKLINE

5MG

N021036 001 Jul 26, 1999

ZANUBRUTINIB

CAPSULE; ORAL

BRUKINSA

+! BEIGENE

80MG

N213217 001 Nov 14, 2019

ZAVEGEPANT HYDROCHLORIDE

SPRAY, METERED; NASAL

ZAVZPRET

+! PFIZER

EQ 10MG BASE/SPRAY

N216386 001 Mar 09, 2023

ZICONOTIDE ACETATE

INJECTABLE; INTRATHECAL

PRIALT

+! TERSERA

100MCG/1ML (100MCG/ML)

N021060 002 Dec 28, 2004

+! 500MCG/20ML (25MCG/ML)

N021060 001 Dec 28, 2004

+! 500MCG/5ML (100MCG/ML)

N021060 004 Dec 28, 2004

ZIDOVUDINE

CAPSULE; ORAL

RETROVIR**AB +! VIIV HLTHCARE****100MG****N019655 001** Mar 19, 1987**ZIDOVUDINE****AB AUROBINDO PHARMA LTD****100MG****A078128 001** Mar 27, 2006**AB CIPLA LTD****100MG****A078349 001** May 23, 2007

INJECTABLE; INJECTION

RETROVIR

+! VIIV HLTHCARE

10MG/ML

N019951 001 Feb 02, 1990

SOLUTION; ORAL

RETROVIR**AA +! VIIV HLTHCARE****50MG/5ML****N019910 001** Sep 28, 1989**ZIDOVUDINE****AA AUROBINDO****50MG/5ML****A077268 001** Sep 19, 2005**AA CIPLA LTD****50MG/5ML****A077981 001** Jun 26, 2008

TABLET; ORAL

ZIDOVUDINE**AB AUROBINDO****300MG****A077267 001** Sep 19, 2005**AB CIPLA****300MG****A090561 001** Oct 27, 2010**AB ! HETERO LABS LTD III****300MG****A090092 001** Apr 25, 2008ZILEUTON

TABLET; ORAL

ZYFLO

+! CHIESI

600MG

N020471 003 Dec 09, 1996

TABLET, EXTENDED RELEASE; ORAL

ZILEUTON**AB AIZANT****600MG****A211390 001** Oct 23, 2020**AB ANNORA PHARMA****600MG****A215742 001** Oct 11, 2022**AB ! RISING****600MG****A204929 001** Mar 17, 2017**AB STRIDES PHARMA****600MG****A212670 001** Dec 16, 2019ZILUCOPLAN SODIUM

SOLUTION; SUBCUTANEOUS

ZILBRYSQ

+! UCB INC

EQ 16.6MG BASE/0.416ML (EQ 16.6MG
BASE/0.416ML)

N216834 001 Oct 17, 2023

+! EQ 23MG BASE/0.574ML (EQ 23MG
BASE/0.574ML)

N216834 002 Oct 17, 2023

+! EQ 32.4MG BASE/0.81ML (EQ 32.4
BASE/0.81ML)

N216834 003 Oct 17, 2023

ZINC ACETATE

CAPSULE; ORAL

GALZIN

+ TEVA

EQ 25MG ZINC

N020458 001 Jan 28, 1997

+! EQ 50MG ZINC

N020458 002 Jan 28, 1997

PRESCRIPTION DRUG PRODUCT LIST

ZINC CHLORIDE

INJECTABLE; INJECTION

ZINC CHLORIDE

<u>AP</u>	EXELA PHARMA	<u>EQ 1MG ZINC/ML</u>	<u>A212007 001</u>	May 21, 2021
<u>AP</u>	SOMERSET	<u>EQ 1MG ZINC/ML</u>	<u>A216152 001</u>	Oct 16, 2024

ZINC CHLORIDE IN PLASTIC CONTAINER

<u>AP</u>	+! HOSPIRA	<u>EQ 1MG ZINC/ML</u>	<u>N018959 001</u>	Jun 26, 1986
-----------	------------	-----------------------	--------------------	--------------

ZINC SULFATE

SOLUTION; INTRAVENOUS

ZINC SULFATE

<u>AP</u>	+! AM REGENT	<u>EQ 10MG BASE/10ML (EQ 1MG BASE/ML)</u>	<u>N209377 003</u>	Apr 15, 2020
<u>AP</u>	+!	<u>EQ 25MG BASE/5ML (EQ 5MG BASE/ML)</u>	<u>N209377 002</u>	Jul 18, 2019
<u>AP</u>	+!	<u>EQ 30MG BASE/10ML (EQ 3MG BASE/ML)</u>	<u>N209377 001</u>	Jul 18, 2019
<u>AP</u>	APOTEX	<u>EQ 10MG BASE/10ML (EQ 1MG BASE/ML)</u>	<u>A218059 002</u>	Oct 28, 2024
<u>AP</u>		<u>EQ 25MG BASE/5ML (EQ 5MG BASE/ML)</u>	<u>A218059 001</u>	Jun 10, 2024
<u>AP</u>	FRESENIUS KABI USA	<u>EQ 10MG BASE/10ML (EQ 1MG BASE/ML)</u>	<u>A216145 001</u>	Dec 27, 2022
<u>AP</u>		<u>EQ 25MG BASE/5ML (EQ 5MG BASE/ML)</u>	<u>A216145 002</u>	Dec 27, 2022
<u>AP</u>		<u>EQ 30MG BASE/10ML (EQ 3MG BASE/ML)</u>	<u>A216145 003</u>	Dec 27, 2022
<u>AP</u>	GLAND PHARMA LTD	<u>EQ 10MG BASE/10ML (EQ 1MG BASE/ML)</u>	<u>A216249 003</u>	Sep 01, 2023
<u>AP</u>		<u>EQ 25MG BASE/5ML (EQ 5MG BASE/ML)</u>	<u>A216249 001</u>	May 03, 2022
<u>AP</u>		<u>EQ 30MG BASE/10ML (EQ 3MG BASE/ML)</u>	<u>A216249 002</u>	May 03, 2022
<u>AP</u>	NIVAGEN PHARMS INC	<u>EQ 10MG BASE/10ML (EQ 1MG BASE/ML)</u>	<u>A214597 001</u>	Jul 05, 2024
<u>AP</u>		<u>EQ 25MG BASE/5ML (EQ 5MG BASE/ML)</u>	<u>A214597 002</u>	Jul 05, 2024
<u>AP</u>		<u>EQ 30MG BASE/10ML (EQ 3MG BASE/ML)</u>	<u>A214597 003</u>	Jul 05, 2024
<u>AP</u>	SOMERSET THERAPS LLC	<u>EQ 25MG BASE/5ML (EQ 5MG BASE/ML)</u>	<u>A216135 001</u>	Jul 17, 2024
<u>AP</u>		<u>EQ 30MG BASE/10ML (EQ 3MG BASE/ML)</u>	<u>A216135 002</u>	Jul 17, 2024
<u>AP</u>	ZYDUS PHARMS	<u>EQ 30MG BASE/10ML (EQ 3MG BASE/ML)</u>	<u>A217074 003</u>	Aug 22, 2023
<u>AP</u>		<u>EQ 10MG BASE/10ML (EQ 1MG BASE/ML)</u>	<u>A217074 001</u>	Aug 22, 2023
<u>AP</u>		<u>EQ 25MG BASE/5ML (EQ 5MG BASE/ML)</u>	<u>A217074 002</u>	Aug 22, 2023

ZIPRASIDONE HYDROCHLORIDE

CAPSULE; ORAL

GEODON

<u>AB</u>	+! VIATRIS	<u>EQ 20MG BASE</u>	<u>N020825 001</u>	Feb 05, 2001
<u>AB</u>	+	<u>EQ 40MG BASE</u>	<u>N020825 002</u>	Feb 05, 2001
<u>AB</u>	+	<u>EQ 60MG BASE</u>	<u>N020825 003</u>	Feb 05, 2001
<u>AB</u>	+	<u>EQ 80MG BASE</u>	<u>N020825 004</u>	Feb 05, 2001

ZIPRASIDONE HYDROCHLORIDE

<u>AB</u>	APOTEX	<u>EQ 20MG BASE</u>	<u>A077561 001</u>	Mar 02, 2012
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A077561 002</u>	Mar 02, 2012
<u>AB</u>		<u>EQ 60MG BASE</u>	<u>A077561 003</u>	Mar 02, 2012
<u>AB</u>		<u>EQ 80MG BASE</u>	<u>A077561 004</u>	Mar 02, 2012
<u>AB</u>	AUROBINDO PHARMA	<u>EQ 20MG BASE</u>	<u>A204117 001</u>	Dec 27, 2016
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A204117 002</u>	Dec 27, 2016
<u>AB</u>		<u>EQ 60MG BASE</u>	<u>A204117 003</u>	Dec 27, 2016
<u>AB</u>		<u>EQ 80MG BASE</u>	<u>A204117 004</u>	Dec 27, 2016
<u>AB</u>	CHARTWELL RX	<u>EQ 20MG BASE</u>	<u>A090348 001</u>	Sep 05, 2012
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A090348 002</u>	Sep 05, 2012
<u>AB</u>		<u>EQ 60MG BASE</u>	<u>A090348 003</u>	Sep 05, 2012
<u>AB</u>		<u>EQ 80MG BASE</u>	<u>A090348 004</u>	Sep 05, 2012
<u>AB</u>	DR REDDYS LABS INC	<u>EQ 20MG BASE</u>	<u>A077565 001</u>	Mar 02, 2012
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A077565 002</u>	Mar 02, 2012
<u>AB</u>		<u>EQ 60MG BASE</u>	<u>A077565 003</u>	Mar 02, 2012
<u>AB</u>		<u>EQ 80MG BASE</u>	<u>A077565 004</u>	Mar 02, 2012
<u>AB</u>	LUPIN PHARMS	<u>EQ 20MG BASE</u>	<u>A077560 001</u>	Mar 02, 2012
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A077560 002</u>	Mar 02, 2012
<u>AB</u>		<u>EQ 60MG BASE</u>	<u>A077560 003</u>	Mar 02, 2012
<u>AB</u>		<u>EQ 80MG BASE</u>	<u>A077560 004</u>	Mar 02, 2012
<u>AB</u>	MACLEODS PHARMS LTD	<u>EQ 20MG BASE</u>	<u>A204375 001</u>	Feb 17, 2017
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A204375 002</u>	Feb 17, 2017
<u>AB</u>		<u>EQ 60MG BASE</u>	<u>A204375 003</u>	Feb 17, 2017
<u>AB</u>		<u>EQ 80MG BASE</u>	<u>A204375 004</u>	Feb 17, 2017
<u>AB</u>	SANDOZ INC	<u>EQ 20MG BASE</u>	<u>A077562 001</u>	Jun 01, 2012
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A077562 002</u>	Jun 01, 2012
<u>AB</u>		<u>EQ 60MG BASE</u>	<u>A077562 003</u>	Jun 01, 2012
<u>AB</u>		<u>EQ 80MG BASE</u>	<u>A077562 004</u>	Jun 01, 2012
<u>AB</u>	ZYDUS LIFESCIENCES	<u>EQ 20MG BASE</u>	<u>A208988 001</u>	Aug 22, 2017
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A208988 002</u>	Aug 22, 2017
<u>AB</u>		<u>EQ 60MG BASE</u>	<u>A208988 003</u>	Aug 22, 2017
<u>AB</u>		<u>EQ 80MG BASE</u>	<u>A208988 004</u>	Aug 22, 2017

PRESCRIPTION DRUG PRODUCT LIST

ZIPRASIDONE MESYLATE

POWDER; INTRAMUSCULAR

GEODON

<u>AP</u>	<u>+!</u>	<u>VIATRIS</u>	<u>EQ 20MG BASE/VIAL</u>	<u>N020919</u>	<u>001</u>	Jun 21, 2002
-----------	-----------	----------------	--------------------------	----------------	------------	--------------

ZIPRASIDONE MESYLATE

<u>AP</u>		<u>GLAND PHARMA LTD</u>	<u>EQ 20MG BASE/VIAL</u>	<u>A211908</u>	<u>001</u>	Dec 26, 2019
-----------	--	-------------------------	--------------------------	----------------	------------	--------------

<u>AP</u>		<u>MSN</u>	<u>EQ 20MG BASE/VIAL</u>	<u>A216091</u>	<u>001</u>	Sep 15, 2022
-----------	--	------------	--------------------------	----------------	------------	--------------

ZOLEDRONIC ACID

INJECTABLE; INTRAVENOUS

RECLAST

<u>AP</u>	<u>+!</u>	<u>SANDOZ</u>	<u>EQ 5MG BASE/100ML</u>	<u>N021817</u>	<u>001</u>	Apr 16, 2007
-----------	-----------	---------------	--------------------------	----------------	------------	--------------

ZOLEDRONIC

<u>AP</u>	<u>!</u>	<u>GLAND PHARMA LTD</u>	<u>EQ 4MG BASE/100ML</u>	<u>A205749</u>	<u>001</u>	Jun 29, 2018
-----------	----------	-------------------------	--------------------------	----------------	------------	--------------

ZOLEDRONIC ACID

<u>AP</u>		<u>ACCORD HLTHCARE</u>	<u>EQ 4MG BASE/5ML</u>	<u>A205279</u>	<u>001</u>	Nov 28, 2016
-----------	--	------------------------	------------------------	----------------	------------	--------------

<u>AP</u>		<u>AMNEAL</u>	<u>EQ 4MG BASE/100ML</u>	<u>A210174</u>	<u>001</u>	Oct 27, 2017
-----------	--	---------------	--------------------------	----------------	------------	--------------

<u>AP</u>		<u>APOTEX</u>	<u>EQ 5MG BASE/100ML</u>	<u>A204367</u>	<u>001</u>	Dec 24, 2015
-----------	--	---------------	--------------------------	----------------	------------	--------------

<u>AP</u>		<u>AVET LIFESCIENCES</u>	<u>EQ 4MG BASE/5ML</u>	<u>A201783</u>	<u>001</u>	Mar 12, 2013
-----------	--	--------------------------	------------------------	----------------	------------	--------------

<u>AP</u>		<u>BPI LABS</u>	<u>EQ 4MG BASE/5ML</u>	<u>A207341</u>	<u>001</u>	Dec 29, 2017
-----------	--	-----------------	------------------------	----------------	------------	--------------

<u>AP</u>		<u>CHARTWELL RX</u>	<u>EQ 4MG BASE/5ML</u>	<u>A202571</u>	<u>001</u>	May 07, 2013
-----------	--	---------------------	------------------------	----------------	------------	--------------

<u>AP</u>			<u>EQ 5MG BASE/100ML</u>	<u>A202163</u>	<u>001</u>	Aug 05, 2013
-----------	--	--	--------------------------	----------------	------------	--------------

<u>AP</u>	<u>!</u>	<u>DR REDDYS LABS LTD</u>	<u>EQ 4MG BASE/5ML</u>	<u>A091186</u>	<u>001</u>	Mar 04, 2013
-----------	----------	---------------------------	------------------------	----------------	------------	--------------

<u>AP</u>			<u>EQ 5MG BASE/100ML</u>	<u>A091363</u>	<u>001</u>	Mar 29, 2013
-----------	--	--	--------------------------	----------------	------------	--------------

<u>AP</u>		<u>EPIC PHARMA LLC</u>	<u>EQ 4MG BASE/5ML</u>	<u>A202548</u>	<u>001</u>	May 22, 2014
-----------	--	------------------------	------------------------	----------------	------------	--------------

<u>AP</u>			<u>EQ 5MG BASE/100ML</u>	<u>A200918</u>	<u>001</u>	Aug 21, 2014
-----------	--	--	--------------------------	----------------	------------	--------------

<u>AP</u>		<u>FRESENIUS KABI USA</u>	<u>EQ 4MG BASE/5ML</u>	<u>A091516</u>	<u>001</u>	Apr 23, 2015
-----------	--	---------------------------	------------------------	----------------	------------	--------------

<u>AP</u>		<u>GLAND</u>	<u>EQ 5MG BASE/100ML</u>	<u>A204217</u>	<u>001</u>	Aug 18, 2016
-----------	--	--------------	--------------------------	----------------	------------	--------------

<u>AP</u>		<u>GLAND PHARMA LTD</u>	<u>EQ 4MG BASE/5ML</u>	<u>A202930</u>	<u>001</u>	Aug 05, 2013
-----------	--	-------------------------	------------------------	----------------	------------	--------------

<u>AP</u>			<u>EQ 5MG BASE/100ML</u>	<u>A209578</u>	<u>001</u>	Aug 08, 2019
-----------	--	--	--------------------------	----------------	------------	--------------

<u>AP</u>		<u>HIKMA FARMACEUTICA</u>	<u>EQ 4MG BASE/5ML</u>	<u>A202182</u>	<u>001</u>	Jun 03, 2013
-----------	--	---------------------------	------------------------	----------------	------------	--------------

<u>AP</u>		<u>HOSPIRA INC</u>	<u>EQ 5MG BASE/100ML</u>	<u>A202837</u>	<u>001</u>	Apr 05, 2013
-----------	--	--------------------	--------------------------	----------------	------------	--------------

<u>AP</u>	<u>+!</u>	<u>INFORLIFE</u>	<u>EQ 4MG BASE/100ML</u>	<u>N203231</u>	<u>001</u>	Aug 02, 2013
-----------	-----------	------------------	--------------------------	----------------	------------	--------------

<u>AP</u>			<u>EQ 5MG BASE/100ML</u>	<u>A202828</u>	<u>001</u>	Sep 23, 2013
-----------	--	--	--------------------------	----------------	------------	--------------

<u>AP</u>		<u>MEITHEAL</u>	<u>EQ 5MG BASE/100ML</u>	<u>A213371</u>	<u>001</u>	Jun 05, 2023
-----------	--	-----------------	--------------------------	----------------	------------	--------------

<u>AP</u>		<u>MYLAN LABS LTD</u>	<u>EQ 4MG BASE/5ML</u>	<u>A202650</u>	<u>001</u>	Mar 04, 2013
-----------	--	-----------------------	------------------------	----------------	------------	--------------

<u>AP</u>			<u>EQ 5MG BASE/100ML</u>	<u>A203841</u>	<u>001</u>	Feb 14, 2017
-----------	--	--	--------------------------	----------------	------------	--------------

<u>AP</u>			<u>EQ 5MG BASE/100ML</u>	<u>A205254</u>	<u>001</u>	Oct 27, 2017
-----------	--	--	--------------------------	----------------	------------	--------------

<u>AP</u>		<u>SAGENT PHARMS INC</u>	<u>EQ 4MG BASE/5ML</u>	<u>A091493</u>	<u>001</u>	Nov 24, 2014
-----------	--	--------------------------	------------------------	----------------	------------	--------------

<u>AP</u>		<u>USV</u>	<u>EQ 4MG BASE/5ML</u>	<u>A202923</u>	<u>001</u>	Sep 04, 2014
-----------	--	------------	------------------------	----------------	------------	--------------

SOLUTION; INTRAVENOUS

ZOLEDRONIC ACID

		<u>HOSPIRA</u>	<u>EQ 4MG BASE/100ML (EQ 0.04MG BASE/ML)</u>	<u>N204016</u>	<u>001</u>	Dec 28, 2015
--	--	----------------	--	----------------	------------	--------------

ZOLMITRIPTAN

SPRAY; NASAL

ZOLMITRIPTAN

<u>AB</u>		<u>PADAGIS ISRAEL</u>	<u>2.5MG/SPRAY</u>	<u>A212469</u>	<u>001</u>	Sep 30, 2021
-----------	--	-----------------------	--------------------	----------------	------------	--------------

<u>AB</u>	<u>!</u>		<u>5MG/SPRAY</u>	<u>A212469</u>	<u>002</u>	Sep 30, 2021
-----------	----------	--	------------------	----------------	------------	--------------

ZOMIG

<u>AB</u>	<u>+</u>	<u>AMNEAL</u>	<u>2.5MG/SPRAY</u>	<u>N021450</u>	<u>003</u>	Sep 16, 2013
-----------	----------	---------------	--------------------	----------------	------------	--------------

<u>AB</u>	<u>+!</u>		<u>5MG/SPRAY</u>	<u>N021450</u>	<u>004</u>	Sep 30, 2003
-----------	-----------	--	------------------	----------------	------------	--------------

TABLET; ORAL

ZOLMITRIPTAN

<u>AB</u>		<u>AJANTA PHARMA LTD</u>	<u>2.5MG</u>	<u>A204041</u>	<u>001</u>	May 20, 2016
-----------	--	--------------------------	--------------	----------------	------------	--------------

<u>AB</u>	<u>!</u>		<u>5MG</u>	<u>A204041</u>	<u>002</u>	May 20, 2016
-----------	----------	--	------------	----------------	------------	--------------

<u>AB</u>		<u>ALEMBIC</u>	<u>2.5MG</u>	<u>A204232</u>	<u>001</u>	Sep 30, 2015
-----------	--	----------------	--------------	----------------	------------	--------------

<u>AB</u>			<u>5MG</u>	<u>A204232</u>	<u>002</u>	Sep 30, 2015
-----------	--	--	------------	----------------	------------	--------------

<u>AB</u>		<u>AUROBINDO PHARMA</u>	<u>2.5MG</u>	<u>A207021</u>	<u>001</u>	May 11, 2016
-----------	--	-------------------------	--------------	----------------	------------	--------------

<u>AB</u>			<u>5MG</u>	<u>A207021</u>	<u>002</u>	May 11, 2016
-----------	--	--	------------	----------------	------------	--------------

<u>AB</u>		<u>GLENMARK PHARMS LTD</u>	<u>2.5MG</u>	<u>A201779</u>	<u>001</u>	May 14, 2013
-----------	--	----------------------------	--------------	----------------	------------	--------------

<u>AB</u>			<u>5MG</u>	<u>A201779</u>	<u>002</u>	May 14, 2013
-----------	--	--	------------	----------------	------------	--------------

<u>AB</u>		<u>JUBILANT GENERICS</u>	<u>2.5MG</u>	<u>A202279</u>	<u>001</u>	Nov 20, 2014
-----------	--	--------------------------	--------------	----------------	------------	--------------

<u>AB</u>			<u>5MG</u>	<u>A202279</u>	<u>002</u>	Nov 20, 2014
-----------	--	--	------------	----------------	------------	--------------

<u>AB</u>		<u>ORBION PHARMS</u>	<u>2.5MG</u>	<u>A203726</u>	<u>001</u>	Oct 21, 2019
-----------	--	----------------------	--------------	----------------	------------	--------------

<u>AB</u>			<u>5MG</u>	<u>A203726</u>	<u>002</u>	Oct 21, 2019
-----------	--	--	------------	----------------	------------	--------------

<u>AB</u>		<u>PLD ACQUISITIONS</u>	<u>2.5MG</u>	<u>A207867</u>	<u>001</u>	Feb 27, 2017
-----------	--	-------------------------	--------------	----------------	------------	--------------

<u>AB</u>		<u>LLC</u>				
-----------	--	------------	--	--	--	--

<u>AB</u>			<u>5MG</u>	<u>A207867</u>	<u>002</u>	Feb 27, 2017
-----------	--	--	------------	----------------	------------	--------------

<u>AB</u>		<u>SOMERSET THERAPS</u>	<u>2.5MG</u>	<u>A206973</u>	<u>001</u>	Jun 30, 2017
-----------	--	-------------------------	--------------	----------------	------------	--------------

<u>AB</u>		<u>LLC</u>				
-----------	--	------------	--	--	--	--

<u>AB</u>			<u>5MG</u>	<u>A206973</u>	<u>002</u>	Jun 30, 2017
-----------	--	--	------------	----------------	------------	--------------

<u>AB</u>		<u>ZYDUS PHARMS</u>	<u>2.5MG</u>	<u>A203019</u>	<u>001</u>	Jul 11, 2018
-----------	--	---------------------	--------------	----------------	------------	--------------

<u>AB</u>			<u>5MG</u>	<u>A203019</u>	<u>002</u>	Jul 11, 2018
-----------	--	--	------------	----------------	------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

ZOLMITRIPTAN

TABLET, ORALLY DISINTEGRATING;ORAL

ZOLMITRIPTAN

<u>AB</u>	ALEMBIC	<u>2.5MG</u>	<u>A205074 001</u>	Dec 01, 2016
<u>AB</u>		<u>5MG</u>	<u>A205074 002</u>	Dec 01, 2016
<u>AB</u>	GLENMARK PHARMS LTD	<u>2.5MG</u>	<u>A202560 001</u>	May 14, 2013
<u>AB</u>		<u>5MG</u>	<u>A202560 002</u>	May 14, 2013
<u>AB</u>	JUBILANT GENERICS	<u>2.5MG</u>	<u>A202956 001</u>	Sep 17, 2015
<u>AB</u>		<u>5MG</u>	<u>A202956 002</u>	Sep 17, 2015
<u>AB</u>	ZYDUS PHARMS USA INC	<u>2.5MG</u>	<u>A202890 001</u>	May 15, 2013
<u>AB</u>	!	<u>5MG</u>	<u>A202890 002</u>	May 15, 2013

ZOLPIDEM TARTRATE

CAPSULE;ORAL

ZOLPIDEM TARTRATE

+! ALMATICA

7.5MG

N215721 001 May 09, 2023

TABLET;ORAL

AMBIEN

<u>AB</u>	+	SANOFI AVENTIS US	<u>5MG</u>	<u>N019908 001</u>	Dec 16, 1992
<u>AB</u>	+	!	<u>10MG</u>	<u>N019908 002</u>	Dec 16, 1992

ZOLPIDEM TARTRATE

<u>AB</u>	ACME LABS	<u>5MG</u>	<u>A077214 001</u>	Apr 23, 2007
<u>AB</u>		<u>10MG</u>	<u>A077214 002</u>	Apr 23, 2007
<u>AB</u>	APOTEX INC	<u>5MG</u>	<u>A077884 001</u>	Apr 23, 2007
<u>AB</u>		<u>10MG</u>	<u>A077884 002</u>	Apr 23, 2007
<u>AB</u>	AUROBINDO PHARMA	<u>5MG</u>	<u>A078413 001</u>	May 04, 2007
<u>AB</u>		<u>10MG</u>	<u>A078413 002</u>	May 04, 2007
<u>AB</u>	CHARTWELL MOLECULAR	<u>5MG</u>	<u>A077388 001</u>	Jul 30, 2012
<u>AB</u>		<u>10MG</u>	<u>A077388 002</u>	Jul 30, 2012
<u>AB</u>	SANDOZ	<u>5MG</u>	<u>A077322 001</u>	Apr 23, 2007
<u>AB</u>		<u>10MG</u>	<u>A077322 002</u>	Apr 23, 2007
<u>AB</u>	TEVA	<u>5MG</u>	<u>A076410 001</u>	Apr 23, 2007
<u>AB</u>		<u>10MG</u>	<u>A076410 002</u>	Apr 23, 2007
<u>AB</u>	TORRENT PHARMS	<u>5MG</u>	<u>A077903 001</u>	Aug 17, 2007
<u>AB</u>		<u>10MG</u>	<u>A077903 002</u>	Aug 17, 2007

TABLET;SUBLINGUAL

EDLUAR

<u>AB</u>	+	MYLAN SPECIALITY LP	<u>5MG</u>	<u>N021997 001</u>	Mar 13, 2009
<u>AB</u>	+	!	<u>10MG</u>	<u>N021997 002</u>	Mar 13, 2009

ZOLPIDEM TARTRATE

<u>AB</u>	DR REDDYS	<u>1.75MG</u>	<u>A204503 001</u>	Nov 18, 2016
<u>AB</u>		<u>3.5MG</u>	<u>A204503 002</u>	Nov 18, 2016
<u>AB</u>	ENDO OPERATIONS	<u>1.75MG</u>	<u>A204229 001</u>	Sep 11, 2017
<u>AB</u>		<u>3.5MG</u>	<u>A204229 002</u>	Sep 11, 2017
<u>AB</u>		<u>5MG</u>	<u>A201509 001</u>	Aug 01, 2016
<u>AB</u>		<u>10MG</u>	<u>A201509 002</u>	Aug 01, 2016
<u>AB</u>	NOVEL LABS INC	<u>1.75MG</u>	<u>A204299 001</u>	Jun 03, 2015
<u>AB</u>	!	<u>3.5MG</u>	<u>A204299 002</u>	Jun 03, 2015

TABLET, EXTENDED RELEASE;ORAL

AMBIEN CR

<u>AB</u>	+	SANOFI AVENTIS US	<u>6.25MG</u>	<u>N021774 002</u>	Sep 02, 2005
<u>AB</u>	+	!	<u>12.5MG</u>	<u>N021774 001</u>	Sep 02, 2005

ZOLPIDEM TARTRATE

<u>AB</u>	APOTEX	<u>6.25MG</u>	<u>A200266 001</u>	Sep 10, 2013
<u>AB</u>		<u>12.5MG</u>	<u>A200266 002</u>	Sep 10, 2013
<u>AB</u>	LUPIN LTD	<u>6.25MG</u>	<u>A078970 001</u>	Sep 11, 2013
<u>AB</u>		<u>12.5MG</u>	<u>A078970 002</u>	Sep 11, 2013
<u>AB</u>	SANDOZ	<u>6.25MG</u>	<u>A090107 001</u>	Jul 01, 2011
<u>AB</u>		<u>12.5MG</u>	<u>A090107 002</u>	Jul 01, 2011
<u>AB</u>	SUN PHARM	<u>6.25MG</u>	<u>A204170 001</u>	Jan 24, 2017
<u>AB</u>		<u>12.5MG</u>	<u>A204170 002</u>	Jan 24, 2017

ZONISAMIDE

CAPSULE;ORAL

ZONEGRAN

<u>AB</u>	+	ADVANSZ PHARMA	<u>25MG</u>	<u>N020789 003</u>	Aug 22, 2003
<u>AB</u>	+	!	<u>100MG</u>	<u>N020789 001</u>	Mar 27, 2000

ZONISAMIDE

<u>AB</u>	APOTEX INC	<u>25MG</u>	<u>A077642 001</u>	Dec 22, 2005
<u>AB</u>		<u>50MG</u>	<u>A077642 002</u>	Dec 22, 2005
<u>AB</u>		<u>100MG</u>	<u>A077642 003</u>	Dec 22, 2005
<u>AB</u>	AUROBINDO PHARMA LTD	<u>25MG</u>	<u>A077645 002</u>	Sep 29, 2006

PRESCRIPTION DRUG PRODUCT LIST

ZONISAMIDE

CAPSULE; ORAL

ZONISAMIDE

<u>AB</u>		<u>50MG</u>	<u>A077645 003</u>	Sep 29, 2006
<u>AB</u>		<u>100MG</u>	<u>A077645 001</u>	Dec 22, 2005
<u>AB</u>	BIONPHARMA	<u>25MG</u>	<u>A077813 001</u>	Aug 16, 2006
<u>AB</u>		<u>50MG</u>	<u>A077813 002</u>	Aug 16, 2006
<u>AB</u>		<u>100MG</u>	<u>A077813 003</u>	Aug 16, 2006
<u>AB</u>	GLENMARK PHARMS INC	<u>25MG</u>	<u>A077651 001</u>	Jan 30, 2006
<u>AB</u>		<u>50MG</u>	<u>A077651 002</u>	Jan 30, 2006
<u>AB</u>		<u>100MG</u>	<u>A077651 003</u>	Jan 30, 2006
<u>AB</u>	GRANULES	<u>25MG</u>	<u>A077636 003</u>	Jul 27, 2006
<u>AB</u>		<u>50MG</u>	<u>A077636 002</u>	Jul 27, 2006
<u>AB</u>		<u>100MG</u>	<u>A077636 001</u>	Dec 22, 2005
<u>AB</u>	INVAGEN PHARMS	<u>25MG</u>	<u>A077869 001</u>	May 31, 2006
<u>AB</u>		<u>50MG</u>	<u>A077869 002</u>	May 31, 2006
<u>AB</u>		<u>100MG</u>	<u>A077869 003</u>	May 31, 2006
<u>AB</u>	SUN PHARM INDS (IN)	<u>25MG</u>	<u>A077634 001</u>	Mar 17, 2006
<u>AB</u>		<u>50MG</u>	<u>A077634 002</u>	Mar 17, 2006
<u>AB</u>		<u>100MG</u>	<u>A077634 003</u>	Mar 17, 2006
<u>AB</u>	UNICHEM	<u>25MG</u>	<u>A214492 001</u>	Jan 26, 2021
<u>AB</u>		<u>50MG</u>	<u>A214492 002</u>	Jan 26, 2021
<u>AB</u>		<u>100MG</u>	<u>A214492 003</u>	Jan 26, 2021
<u>AB</u>	ZYDUS LIFESCIENCES	<u>25MG</u>	<u>A077625 001</u>	Oct 16, 2006
<u>AB</u>		<u>50MG</u>	<u>A077625 002</u>	Oct 16, 2006
<u>AB</u>		<u>100MG</u>	<u>A077625 003</u>	Oct 16, 2006

SUSPENSION; ORAL

ZONISADE

+! AZURITY

100MG/5ML

N214273 001 Jul 15, 2022

ZURANOLONE

CAPSULE; ORAL

ZURZUVAE

+ BIOGEN INC

20MG

N217369 001 Oct 31, 2023

+

25MG

N217369 002 Oct 31, 2023

+!

30MG

N217369 003 Oct 31, 2023

OTC DRUG PRODUCT LIST

ACETAMINOPHEN

SUPPOSITORY; RECTAL

ACETAMINOPHEN

PERRIGO NEW YORK	120MG	A070607	001	Apr 06, 1987
	650MG	A070608	001	Dec 01, 1986
+ TARO	120MG	N018337	003	Sep 12, 1983
+	325MG	N018337	002	

INFANTS' FEVERALL

+ TARO	80MG	N018337	004	Aug 26, 1992
--------	------	---------	-----	--------------

TABLET, EXTENDED RELEASE; ORAL

ACETAMINOPHEN

AUROBINDO PHARMA	650MG	A207229	001	Nov 09, 2016
GRANULES	650MG	A211544	001	Apr 16, 2019
HERITAGE PHARMA	650MG	A207035	001	May 31, 2018
MARKSANS PHARMA	650MG	A215486	001	Aug 25, 2021
OHM LABS	650MG	A076200	001	Mar 19, 2002
PERRIGO	650MG	A075077	001	Feb 25, 2000

TYLENOL

+! KENVUE BRANDS	650MG	N019872	001	Jun 08, 1994
+!	650MG	N019872	002	Jan 11, 2001

ACETAMINOPHEN; ASPIRIN; CAFFEINE

TABLET; ORAL

ACETAMINOPHEN, ASPIRIN AND CAFFEINE

AUROBINDO PHARMA	250MG; 250MG; 65MG	A211695	001	Feb 02, 2022
LTD				
GRANULES	250MG; 250MG; 65MG	A214039	001	Feb 23, 2021
PERRIGO	250MG; 250MG; 65MG	A075794	001	Nov 26, 2001

EXCEDRIN (MIGRAINE RELIEF)

+! HALEON US HOLDINGS	250MG; 250MG; 65MG	N020802	001	Jan 14, 1998
-----------------------	--------------------	---------	-----	--------------

ACETAMINOPHEN; IBUPROFEN

TABLET; ORAL

ACETAMINOPHEN AND IBUPROFEN

AUROBINDO PHARMA	250MG; 125MG	A218359	001	Mar 26, 2024
LTD				
BIONPHARMA	250MG; 125MG	A216999	001	Aug 01, 2023
DR REDDYS	250MG; 125MG	A218247	001	Aug 23, 2024
GLENMARK PHARMS LTD	250MG; 125MG	A218311	001	Apr 26, 2024
GRANULES	250MG; 125MG	A216592	001	Jul 13, 2023
L PERRIGO CO	250MG; 125MG	A214836	001	Feb 28, 2023
MARKSANS PHARMA	250MG; 125MG	A216994	001	Jul 10, 2023

ADVIL DUAL ACTION WITH ACETAMINOPHEN

+! HALEON US HOLDINGS	250MG; 125MG	N211733	001	Feb 28, 2020
-----------------------	--------------	---------	-----	--------------

ADAPALENE

GEL; TOPICAL

ADAPALENE

GLENMARK PHARMS INC	0.1%	A091314	001	Jul 01, 2010
P AND L	0.1%	A090962	001	Jun 02, 2010
TARO	0.1%	A215940	001	Jan 14, 2022

DIFFERIN

+! GALDERMA LABS LP	0.1%	N020380	002	Jul 08, 2016
---------------------	------	---------	-----	--------------

ALCAFTADINE

SOLUTION/DROPS; OPHTHALMIC

ALCAFTADINE

ALEMBIC	0.25%	A209290	001	Oct 02, 2024
GLAND PHARMA LTD	0.25%	A209706	001	Mar 01, 2024

LASTACFT

+! ABBVIE	0.25%	N022134	001	Jul 28, 2010
-----------	-------	---------	-----	--------------

ALCOHOL; CHLORHEXIDINE GLUCONATE

SOLUTION; TOPICAL

AVAGARD

+! 3M	61%; 1%	N021074	001	Jun 07, 2001
-------	---------	---------	-----	--------------

ASPIRIN

CAPSULE; ORAL

VAZALORE

+ PLX PHARMA	81MG	N203697	002	Feb 26, 2021
+!	325MG	N203697	001	Jan 14, 2013

OTC DRUG PRODUCT LISTAVOBENZONE; ECAMSULE; OCTOCRYLENE

CREAM; TOPICAL

ANTHELIOS SX

+! LOREAL USA 2%;2%;10%

N021502 001 Jul 21, 2006

CAPITAL SOLEIL 15

+! LOREAL USA 2%;3%;10%

N021501 001 Oct 02, 2006

AVOBENZONE; ECAMSULE; OCTOCRYLENE; TITANIUM DIOXIDE

CREAM; TOPICAL

ANTHELIOS 20

+! LOREAL USA 2%;2%;10%;2%

N021471 001 Oct 05, 2006

ANTHELIOS 40

+! LOREAL USA 2%;3%;10%;5%

N022009 001 Mar 31, 2008

+! 2%;3%;10%;5%

N022009 002 Oct 29, 2009

AZELASTINE HYDROCHLORIDE

SPRAY, METERED; NASAL

ASTEPRO ALLERGY

+! BAYER HLTHCARE 0.2055MG/SPRAY

N213872 001 Jun 17, 2021

AZELASTINE HYDROCHLORIDE

AMNEAL 0.2055MG/SPRAY

A216576 001 Jun 05, 2024

AZELASTINE HYDROCHLORIDE ALLERGY

APOTEX 0.2055MG/SPRAY

A216421 001 May 29, 2024

AZELASTINE HYDROCHLORIDE CHILDREN'S ALLERGY

APOTEX 0.2055MG/SPRAY

A216421 002 May 29, 2024

CHILDREN'S ASTEPRO ALLERGY

+! BAYER HLTHCARE 0.2055MG/SPRAY

N213872 002 Jun 17, 2021

BENTOQUATAM

LOTION; TOPICAL

IVY BLOCK

+! STAND HOMEOPATH 5%

N020532 001 Aug 26, 1996

BRIMONIDINE TARTRATE

SOLUTION/DROPS; OPHTHALMIC

BRIMONIDINE TARTRATE

DR REDDYS LABS SA 0.025%

A216361 001 Feb 16, 2024

LUMIFY

+! BAUSCH AND LOMB INC 0.025%

N208144 001 Dec 22, 2017

LUMIFY PRESERVATIVE FREE

+! BAUSCH AND LOMB INC 0.025%

N218424 001 Apr 19, 2024

BUDESONIDE

SPRAY, METERED; NASAL

BUDESONIDE

APOTEX INC 0.032MG/SPRAY

A078949 002 Nov 20, 2015

RHINOCORT ALLERGY

+! KENVUE BRANDS 0.032MG/SPRAY

N020746 003 Mar 23, 2015

BUTENAFINE HYDROCHLORIDE

CREAM; TOPICAL

BUTENAFINE HYDROCHLORIDE

TARO 1%

A205181 001 Nov 16, 2017

LOTRIMIN ULTRA

+! BAYER HEALTHCARE 1%

N021307 001 Dec 07, 2001

LLC

CALCIUM CARBONATE; FAMOTIDINE; MAGNESIUM HYDROXIDE

TABLET, CHEWABLE; ORAL

FAMOTIDINE, CALCIUM CARBONATE, AND MAGNESIUM HYDROXIDE

PERRIGO R AND D 800MG;10MG;165MG

A077355 001 Feb 06, 2008

PEPCID COMPLETE

+! KENVUE BRANDS 800MG;10MG;165MG

N020958 001 Oct 16, 2000

CETIRIZINE HYDROCHLORIDE

CAPSULE; ORAL

CETIRIZINE HYDROCHLORIDE ALLERGY

AUROBINDO PHARMA 10MG

A209107 001 Jul 20, 2018

LTD

+ BIONPHARMA 5MG

N022429 001 Jul 23, 2009

+! 10MG

N022429 004 Jul 23, 2009

CATALENT 10MG

A213105 001 Sep 21, 2020

STRIDES SOFTGELS 10MG

A205291 001 Jul 21, 2017

CETIRIZINE HYDROCHLORIDE HIVES RELIEF

AUROBINDO PHARMA 10MG

A209107 002 Jul 20, 2018

LTD

+ BIONPHARMA 5MG

N022429 003 Jul 23, 2009

OTC DRUG PRODUCT LIST

CETIRIZINE HYDROCHLORIDE

CAPSULE;ORAL

CETIRIZINE HYDROCHLORIDE HIVES RELIEF

+! 10MG N022429 002 Jul 23, 2009

SOLUTION;ORAL

CHILDREN'S CETIRIZINE HYDROCHLORIDE ALLERGY

AMNEAL PHARMS 5MG/5ML A090765 002 Oct 07, 2009

APOZEAL PHARMS 5MG/5ML A090474 002 Mar 30, 2009

AUROBINDO PHARMA 5MG/5ML A090750 002 Feb 02, 2010

BAJAJ 5MG/5ML A091327 001 Oct 17, 2011

CHARTWELL MOLECULAR 5MG/5ML A091130 001 Apr 22, 2011

HETERO LABS LTD III 5MG/5ML A210622 001 Mar 13, 2019

PERRIGO R AND D 5MG/5ML A204226 001 Sep 09, 2013

5MG/5ML A090254 002 Apr 09, 2008

QUAGEN 5MG/5ML A212266 001 May 16, 2019

TARO 5MG/5ML A090182 002 Apr 22, 2008

5MG/5ML A201546 001 May 20, 2011

CHILDREN'S CETIRIZINE HYDROCHLORIDE HIVES RELIEF

AMNEAL PHARMS 5MG/5ML A090765 001 Oct 07, 2009

APOZEAL PHARMS 5MG/5ML A090474 001 Mar 30, 2009

AUROBINDO PHARMA 5MG/5ML A090750 001 Feb 02, 2010

BAJAJ 5MG/5ML A091327 002 Oct 17, 2011

CHARTWELL MOLECULAR 5MG/5ML A091130 002 Apr 22, 2011

PERRIGO R AND D 5MG/5ML A090254 001 Apr 09, 2008

TARO 5MG/5ML A090182 001 Apr 22, 2008

5MG/5ML A201546 002 May 20, 2011

CHILDREN'S ZYRTEC ALLERGY

+! KENVUE BRANDS 5MG/5ML N022155 002 Nov 16, 2007

CHILDREN'S ZYRTEC HIVES

+! KENVUE BRANDS 5MG/5ML N022155 001 Nov 16, 2007

TABLET;ORAL

CETIRIZINE HYDROCHLORIDE

INDOCO 10MG A218895 001 Oct 03, 2024

CETIRIZINE HYDROCHLORIDE ALLERGY

AMNEAL PHARMS NY 5MG A078780 001 Jan 21, 2010

10MG A078780 004 Jan 21, 2010

APOTEX INC 5MG A078317 001 Dec 27, 2007

10MG A078317 002 Dec 27, 2007

AUROBINDO PHARMA 5MG A090760 001 Aug 05, 2015

LTD 10MG A090760 003 Aug 05, 2015

CONTRACT PHARMACAL 5MG A076047 001 Dec 27, 2007

10MG A076047 002 Dec 27, 2007

DR REDDYS LABS LTD 5MG A078343 004 Jan 15, 2008

10MG A078343 003 Jan 15, 2008

GLENMARK PHARMS INC 5MG A078427 003 Dec 28, 2007

10MG A078427 004 Dec 28, 2007

GRANULES 10MG A209274 001 Dec 22, 2017

MARKSANS PHARMA 5MG A078933 001 Jun 15, 2010

10MG A078933 002 Jun 15, 2010

MYLAN 5MG A076677 001 Dec 27, 2007

10MG A076677 002 Dec 27, 2007

ORBION PHARMS 5MG A078862 001 Feb 19, 2009

10MG A078862 002 Feb 19, 2009

PERRIGO R AND D 5MG A078336 001 Dec 27, 2007

10MG A078336 002 Dec 27, 2007

PLD ACQUISITIONS 5MG A077946 001 Dec 27, 2007

10MG A077946 002 Dec 27, 2007

SUN PHARM INDS LTD 5MG A077498 001 Dec 27, 2007

10MG A077498 002 Dec 27, 2007

TARO 5MG A078072 001 Jul 22, 2009

5MG A078072 003 Jul 22, 2009

UNIQUE 5MG A077829 001 Aug 26, 2009

10MG A077829 004 Aug 26, 2009

CETIRIZINE HYDROCHLORIDE HIVES

DR REDDYS LABS LTD 5MG A078343 001 Jan 15, 2008

10MG A078343 002 Jan 15, 2008

MARKSANS PHARMA 5MG A078933 003 Jun 15, 2010

10MG A078933 004 Jun 15, 2010

MYLAN 5MG A076677 004 Dec 27, 2007

10MG A076677 003 Dec 27, 2007

ORBION PHARMS 5MG A078862 003 Feb 19, 2009

10MG A078862 004 Feb 19, 2009

OTC DRUG PRODUCT LIST

CETIRIZINE HYDROCHLORIDE

TABLET;ORAL

CETIRIZINE HYDROCHLORIDE HIVES

PERRIGO R AND D	5MG	A078336	003	Dec 27, 2007
	10MG	A078336	004	Dec 27, 2007
SUN PHARM INDS LTD	5MG	A077498	003	Dec 27, 2007
	10MG	A077498	004	Dec 27, 2007
UNIQUE	5MG	A077829	003	Aug 26, 2009
!	10MG	A077829	002	Aug 26, 2009

CETIRIZINE HYDROCHLORIDE HIVES RELIEF

AMNEAL PHARMS NY	5MG	A078780	003	Jan 21, 2010
	10MG	A078780	002	Jan 21, 2010
AUROBINDO PHARMA LTD	5MG	A090760	002	Aug 05, 2015
	10MG	A090760	004	Aug 05, 2015
TARO	10MG	A078072	002	Jul 22, 2009
	10MG	A078072	004	Jul 22, 2009

ZYRTEC ALLERGY

+! KENVUE BRANDS	10MG	N019835	004	Nov 16, 2007
------------------	------	---------	-----	--------------

TABLET, CHEWABLE;ORAL

CHILDREN'S CETIRIZINE HYDROCHLORIDE ALLERGY

JUBILANT GENERICS	5MG	A091116	001	Feb 19, 2015
	10MG	A091116	002	Feb 19, 2015
NOVEL LABS INC	5MG	A206793	001	Mar 08, 2016
	10MG	A206793	002	Mar 08, 2016
SANDOZ	5MG	A078692	001	Feb 14, 2008
!	10MG	A078692	002	Feb 14, 2008

CHILDREN'S CETIRIZINE HYDROCHLORIDE HIVES RELIEF

JUBILANT GENERICS	5MG	A091116	003	Feb 19, 2015
	10MG	A091116	004	Feb 19, 2015

CHILDREN'S ZYRTEC ALLERGY

+ KENVUE BRANDS	2.5MG	N021621	007	Nov 30, 2020
+ KENVUE BRANDS	10MG	N021621	004	Nov 16, 2007

TABLET, ORALLY DISINTEGRATING;ORAL

CETIRIZINE HYDROCHLORIDE

AUROBINDO PHARMA LTD	10MG	A213557	001	Sep 11, 2020
----------------------	------	---------	-----	--------------

CETIRIZINE HYDROCHLORIDE ALLERGY

PERRIGO R AND D	10MG	A205490	001	Sep 02, 2015
-----------------	------	---------	-----	--------------

ZYRTEC ALLERGY

+! KENVUE BRANDS	10MG	N022578	001	Sep 03, 2010
------------------	------	---------	-----	--------------

CETIRIZINE HYDROCHLORIDE; PSEUDOEPHEDRINE HYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL

CETIRIZINE HYDROCHLORIDE AND PSEUDOEPHEDRINE HYDROCHLORIDE

IVAX SUB TEVA PHARMS	5MG;120MG	A077170	001	Feb 25, 2008
PLD ACQUISITIONS	5MG;120MG	A077991	001	Mar 05, 2008
PPI-DAC	5MG;120MG	A210719	001	Nov 16, 2018
SUN PHARM INDS LTD	5MG;120MG	A090922	001	Sep 28, 2012
UNICHEM	5MG;120MG	A210507	001	Sep 10, 2024
ZYRTEC-D 12 HOUR				
+! KENVUE BRANDS	5MG;120MG	N021150	002	Nov 09, 2007

CHLORHEXIDINE GLUCONATE

AEROSOL, METERED;TOPICAL

EXIDINE

+! XTTRIUM	4%	N019127	001	Dec 24, 1984
------------	----	---------	-----	--------------

CLOTH;TOPICAL

CHLORHEXIDINE GLUCONATE

+! SAGE PRODS	2%	N021669	001	Apr 25, 2005
---------------	----	---------	-----	--------------

READYPREP CHG

MEDLINE INDUSTRIES	2%	N207964	001	Nov 20, 2018
--------------------	----	---------	-----	--------------

SOLUTION;TOPICAL

CHG SCRUB

ECOLAB	4%	N019258	002	Jul 22, 1986
--------	----	---------	-----	--------------

CHLORHEXIDINE GLUCONATE

BAJAJ	0.75%	N020111	001	Sep 11, 1997
-------	-------	---------	-----	--------------

CIDA-STAT

ECOLAB	2%	N019258	001	Jul 22, 1986
--------	----	---------	-----	--------------

EXIDINE

+! XTTRIUM	2%	N019422	001	Dec 17, 1985
	4%	N019125	001	Dec 24, 1984

OTC DRUG PRODUCT LIST

CHLORHEXIDINE GLUCONATE

SOLUTION;TOPICAL

HIBICLENS

+! MOLNLYCKE HLTH 4% N017768 001

HIBISTAT

+! MOLNLYCKE HLTH 0.5% N018300 001

SPONGE;TOPICAL

BIOSCRUB

GRIFFEN 4% N019822 001 Mar 31, 1989

CHLORHEXIDINE GLUCONATE

! BECTON DICKINSON 4% A072525 001 Oct 24, 1989

CHLORHEXIDINE GLUCONATE; ISOPROPYL ALCOHOL

SOLUTION;TOPICAL

SOLUPREP S

+! 3M HEALTH CARE 2%;70% N208288 001 Aug 08, 2018

SPONGE;TOPICAL

CHLORAPREP ONE-STEP

+! BECTON DICKINSON CO 2%;70% (3ML) N020832 001 Jul 14, 2000

+! 2%;70% (10.5ML) N020832 004 Aug 20, 2003

+! 2%;70% (26ML) N020832 006 Nov 21, 2006

+! 2%;70% (1ML) N020832 008 Oct 23, 2008

CHLORAPREP ONE-STEP FREPP

+! BECTON DICKINSON CO 2%;70% (1.5ML) N020832 003 Apr 26, 2002

CHLORAPREP WITH TINT

+! BECTON DICKINSON CO 2%;70% (26ML) N020832 002 May 03, 2005

+! 2%;70% (10.5ML) N020832 005 Apr 03, 2006

+! 2%;70% (3ML) N020832 007 Oct 10, 2006

SWAB;TOPICAL

CHLORAPREP ONE-STEP SEPP

+! BECTON DICKINSON CO 2%;70% (0.67ML) N021555 001 Oct 07, 2002

CHLORAPREP SINGLE SWABSTICK

+! BECTON DICKINSON CO 2%;70% (1.75ML) N021555 002 May 10, 2005

CHLORAPREP TRIPLE SWABSTICK

+! BECTON DICKINSON CO 2%;70% (5.25ML) N021555 003 Jun 10, 2009

PREVANTICS MAXI SWABSTICK

+! PROF DSPLS 3.15%;70% (5.1ML) N021524 003 Jun 03, 2005

PREVANTICS SWAB

+! PROF DSPLS 3.15%;70% (1ML) N021524 001 Jun 03, 2005

PREVANTICS SWABSTICK

+! PROF DSPLS 3.15%;70% (1.6ML) N021524 002 Jun 03, 2005

CHLORPHENIRAMINE MALEATE

TABLET, EXTENDED RELEASE;ORAL

CHLORPHENIRAMINE MALEATE

! AVANTHI INC 12MG A040829 001 May 13, 2009

CHLORPHENIRAMINE MALEATE; IBUPROFEN; PHENYLEPHRINE HYDROCHLORIDE

TABLET;ORAL

ADVIL ALLERGY AND CONGESTION RELIEF

+! HALEON US HOLDINGS 4MG;200MG;10MG N022113 001 Dec 21, 2011

ADVIL MULTI-SYMP TOM COLD & FLU

+! HALEON US HOLDINGS 4MG;200MG;10MG N022113 002 Apr 28, 2017

CHLORPHENIRAMINE MALEATE; IBUPROFEN; PSEUDOEPHEDRINE HYDROCHLORIDE

SUSPENSION;ORAL

CHILDREN'S ADVIL ALLERGY SINUS

+! HALEON US HOLDINGS 1MG/5ML;100MG/5ML;15MG/5ML N021587 001 Feb 24, 2004

TABLET;ORAL

ADVIL ALLERGY SINUS

+! HALEON US HOLDINGS 2MG;200MG;30MG N021441 001 Dec 19, 2002

CIMETIDINE

TABLET;ORAL

CIMETIDINE

APOTEX 100MG A074948 001 Jun 19, 1998

200MG A074948 002 Jul 26, 2002

L PERRIGO CO 200MG A075285 001 Oct 29, 1998

TAGAMET HB

+! MEDTECH PRODUCTS 200MG N020238 002 Aug 21, 1996

OTC DRUG PRODUCT LIST

CLOTRIMAZOLE

CREAM; VAGINAL

CLOTRIMAZOLE

! P AND L

1%

A074165 001 Jul 16, 1993

TARO

1%

A072641 001 Dec 04, 1995

TRIVAGIZOLE 3

TARO

2%

N021143 001 Apr 12, 2000

CROMOLYN SODIUM

SPRAY, METERED; NASAL

CROMOLYN SODIUM

! BAUSCH AND LOMB

5.2MG/SPRAY

A075702 001 Jul 03, 2001

DEXTROMETHORPHAN HYDROBROMIDE; GUAIFENESIN

TABLET, EXTENDED RELEASE; ORAL

GUAIFENESIN AND DEXTROMETHORPHAN HYDROBROMIDE

ACTAVIS LABS FL

30MG; 600MG

A091070 001 Aug 31, 2015

60MG; 1.2GM

A091070 002 Aug 31, 2015

AMNEAL PHARMS

30MG; 600MG

A209692 001 Nov 01, 2018

60MG; 1.2GM

A209692 002 Nov 01, 2018

AUROBINDO PHARMA

30MG; 600MG

A206941 001 Mar 17, 2017

60MG; 1.2GM

A206941 002 Mar 17, 2017

DR REDDYS

30MG; 600MG

A217340 001 Aug 01, 2023

60MG; 1.2GM

A217340 002 Aug 01, 2023

PERRIGO R AND D

30MG; 600MG

A207602 002 Mar 05, 2018

60MG; 1.2GM

A207602 001 Mar 05, 2018

SUN PHARM

30MG; 600MG

A214781 001 Jul 01, 2021

60MG; 1.2GM

A214781 002 Jul 01, 2021

MUCINEX DM

+ RB HLTH

30MG; 600MG

N021620 002 Apr 29, 2004

+!

60MG; 1.2GM

N021620 001 Apr 29, 2004

DEXTROMETHORPHAN POLISTIREX

SUSPENSION, EXTENDED RELEASE; ORAL

DELSYM

+! RB HLTH

EQ 30MG HYDROBROMIDE/5ML

N018658 001 Oct 08, 1982

DEXTROMETHORPHAN POLISTIREX

AMNEAL

EQ 30MG HYDROBROMIDE/5ML

A203133 001 Jul 28, 2017

TRIS PHARMA INC

EQ 30MG HYDROBROMIDE/5ML

A091135 001 May 25, 2012

DICLOFENAC SODIUM

GEL; TOPICAL

DICLOFENAC SODIUM

AMNEAL PHARMS

1%

A208077 001 Mar 18, 2016

AUROLIFE PHARMA LLC

1%

A204306 001 May 06, 2019

CIPLA

1%

A209903 001 Aug 03, 2018

ENCUBE

1%

A210986 001 Jan 27, 2020

PERRIGO PHARMA INTL

1%

A211253 001 May 16, 2019

VOLTAREN ARTHRITIS PAIN

+! HALEON US HOLDINGS

1%

N022122 001 Oct 17, 2007

DIPHENHYDRAMINE CITRATE; IBUPROFEN

TABLET; ORAL

ADVIL PM

+! HALEON US HOLDINGS

38MG; 200MG

N021394 001 Dec 21, 2005

IBUPROFEN AND DIPHENHYDRAMINE CITRATE

AUROBINDO PHARMA

38MG; 200MG

A216204 001 May 31, 2022

DR REDDYS LABS LTD

38MG; 200MG

A090619 001 Jul 08, 2009

PERRIGO R AND D

38MG; 200MG

A079113 001 Dec 22, 2008

PLD ACQUISITIONS

38MG; 200MG

A211404 001 Apr 11, 2024

LLC

DIPHENHYDRAMINE HYDROCHLORIDE; IBUPROFEN

CAPSULE; ORAL

ADVIL PM

+! HALEON US HOLDINGS

25MG; EQ 200MG FREE ACID AND POTASSIUM SALT

N021393 001 Dec 21, 2005

IBUPROFEN AND DIPHENHYDRAMINE HYDROCHLORIDE

AUROBINDO PHARMA

25MG; EQ 200MG FREE ACID AND POTASSIUM SALT

A210676 001 Feb 14, 2019

LTD

BIONPHARMA

25MG; EQ 200MG FREE ACID AND POTASSIUM SALT

A090397 001 Nov 22, 2010

STRIDES SOFTGELS

25MG; EQ 200MG FREE ACID AND POTASSIUM SALT

A200888 001 Mar 05, 2012

OTC DRUG PRODUCT LIST

DIPHENHYDRAMINE HYDROCHLORIDE; NAPROXEN SODIUM

TABLET; ORAL

ALEVE PM

+! BAYER HLTHCARE 25MG;220MG N205352 001 Jan 17, 2014

NAPROXEN SODIUM AND DIPHENHYDRAMINE HYDROCHLORIDE

AMNEAL PHARMS CO 25MG;220MG A209726 001 Oct 23, 2018

COREPHARMA 25MG;220MG A211830 001 Aug 22, 2019

GRANULES 25MG;220MG A213663 001 Sep 24, 2020

PERRIGO R AND D 25MG;220MG A208499 001 May 10, 2019

DOCOSANOL

CREAM; TOPICAL

ABREVA

+! HALEON US HOLDINGS 10% N020941 001 Jul 25, 2000

DOCOSANOL

ALEMBIC 10% A215839 001 May 03, 2022

AUROBINDO PHARMA 10% A217090 001 Mar 01, 2024

LTD

P AND L 10% A208754 001 Nov 19, 2018

P AND L DEVELOPMENT 10% A212385 001 Oct 07, 2022

TARO 10% A214454 001 Oct 26, 2023

DOXYLAMINE SUCCINATE

TABLET; ORAL

DOXYLAMINE SUCCINATE

LNK 25MG A040564 001 Aug 27, 2004

PERRIGO 25MG A040167 001 Sep 18, 1996

UNISOM

+! CHATTEM 25MG N018066 001

EPINEPHRINE

AEROSOL, METERED; INHALATION

PRIMATENE MIST

+! ARMSTRONG PHARMS 0.125MG/INH N205920 001 Nov 07, 2018

ESOMEPRAZOLE MAGNESIUM

CAPSULE, DELAYED RELEASE; ORAL

ESOMEPRAZOLE MAGNESIUM

AMNEAL PHARMS NY EQ 20MG BASE A209716 001 Jun 05, 2019

ANDA REPOSITORY EQ 20MG BASE A216149 001 Mar 15, 2023

AUROBINDO PHARMA EQ 20MG BASE A209339 001 Oct 16, 2017

DR REDDYS EQ 20MG BASE A207198 001 Apr 30, 2024

EQ 20MG BASE A207673 001 May 15, 2018

GLENMARK SPECLT EQ 20MG BASE A218092 001 Jun 11, 2024

GRAVITI PHARMS EQ 20MG BASE A216349 001 Jun 24, 2022

HETERO LABS LTD III EQ 20MG BASE A212507 001 Jun 02, 2020

MARKSANS PHARMA EQ 20MG BASE A217264 001 Sep 29, 2023

PERRIGO R AND D EQ 20MG BASE A207193 001 Aug 18, 2017

SUN PHARM EQ 20MG BASE A212866 001 May 04, 2020

NEXIUM 24HR

+! ASTRAZENECA LP EQ 20MG BASE N204655 001 Mar 28, 2014

TABLET, DELAYED RELEASE; ORAL

ESOMEPRAZOLE MAGNESIUM

AUROBINDO PHARMA EQ 20MG BASE A214473 001 Jul 12, 2023

LTD

DR REDDYS EQ 20MG BASE A211571 001 May 14, 2020

MYLAN EQ 20MG BASE A212088 001 Jun 25, 2020

NEXIUM 24HR

+! ASTRAZENECA LP EQ 20MG BASE N207920 001 Nov 23, 2015

FAMOTIDINE

TABLET; ORAL

FAMOTIDINE

ANNORA PHARMA 10MG A215766 001 Nov 08, 2021

20MG A215766 002 Nov 08, 2021

ASCENT PHARMS INC 10MG A216030 001 Nov 03, 2021

20MG A216030 002 Nov 03, 2021

AUROBINDO PHARMA 10MG A206531 001 Apr 26, 2016

20MG A206531 002 Apr 26, 2016

CONTRACT PHARMACAL 10MG A218003 001 May 28, 2024

20MG A218003 002 May 28, 2024

DR REDDYS LABS LTD 10MG A077367 002 Aug 17, 2001

20MG A077367 001 Sep 25, 2006

GLENMARK PHARMS INC 10MG A077146 001 Mar 07, 2005

20MG A090837 001 Aug 04, 2010

OTC DRUG PRODUCT LIST

FAMOTIDINE

TABLET; ORAL

FAMOTIDINE

GRAVITI PHARMS	10MG	A218461 001	Mar 14, 2024
	20MG	A218461 002	Mar 14, 2024
MARKSANS PHARMA	10MG	A217543 001	Mar 08, 2023
	20MG	A217543 002	Mar 08, 2023
P AND L	10MG	A075512 001	Jul 26, 2001
PERRIGO	10MG	A075400 001	Mar 18, 2005
PERRIGO R AND D	20MG	A077351 001	Sep 25, 2006
VKT PHARMA	10MG	A215822 001	Jan 28, 2022
	20MG	A215822 002	Jan 28, 2022

PEPCID AC

J AND J CONSUMER INC	10MG	N020902 001	Aug 05, 1999
+ KENVUE BRANDS	10MG	N020325 001	Apr 28, 1995
+!	20MG	N020325 002	Sep 23, 2003

TABLET, CHEWABLE; ORAL

PEPCID AC

+! KENVUE BRANDS	20MG	N020801 002	Dec 17, 2007
------------------	------	-------------	--------------

FEXOFENADINE HYDROCHLORIDE

SUSPENSION; ORAL

CHILDREN'S ALLEGRA ALLERGY

+! CHATTEM SANOFI	30MG/5ML	N201373 001	Jan 24, 2011
-------------------	----------	-------------	--------------

CHILDREN'S FEXOFENADINE HYDROCHLORIDE ALLERGY

P AND L	30MG/5ML	A203330 001	Nov 18, 2014
TARO	30MG/5ML	A208123 001	Nov 09, 2017

CHILDREN'S FEXOFENADINE HYDROCHLORIDE HIVES

! P AND L	30MG/5ML	A203330 002	Nov 18, 2014
TARO	30MG/5ML	A208123 002	Nov 09, 2017

FEXOFENADINE HYDROCHLORIDE

AUROBINDO PHARMA LTD	30MG/5ML	A213466 001	May 23, 2023
-------------------------	----------	-------------	--------------

TABLET; ORAL

ALLEGRA ALLERGY

+ CHATTEM SANOFI	60MG	N020872 007	Jan 24, 2011
+!	180MG	N020872 010	Jan 24, 2011

ALLEGRA HIVES

+! CHATTEM SANOFI	180MG	N020872 009	Jan 24, 2011
-------------------	-------	-------------	--------------

CHILDREN'S FEXOFENADINE HYDROCHLORIDE ALLERGY

AUROBINDO PHARMA LTD	30MG	A202039 001	Nov 19, 2014
DR REDDYS LABS LTD	30MG	A076502 004	Apr 12, 2011
HETERO LABS LTD V	30MG	A204097 001	Aug 19, 2016
TEVA	30MG	A076447 004	Apr 13, 2011
WOCKHARDT	30MG	A079112 002	Feb 08, 2012

CHILDREN'S FEXOFENADINE HYDROCHLORIDE HIVES

DR REDDYS LABS LTD	30MG	A076502 005	Apr 12, 2011
TEVA	30MG	A076447 005	Apr 13, 2011
WOCKHARDT	30MG	A079112 001	Feb 08, 2012

FEXOFENADINE HYDROCHLORIDE

L PERRIGO CO	60MG	A212971 001	Feb 24, 2020
	180MG	A212971 002	Feb 24, 2020

FEXOFENADINE HYDROCHLORIDE ALLERGY

AUROBINDO PHARMA LTD	60MG	A202039 002	Nov 19, 2014
	180MG	A202039 003	Nov 19, 2014
DR REDDYS LABS LTD	60MG	A076502 006	Apr 12, 2011
	180MG	A076502 008	Apr 12, 2011
GRANULES	60MG	A211075 001	Oct 18, 2019
	180MG	A211075 002	Oct 18, 2019
HETERO LABS LTD V	60MG	A204097 002	Aug 19, 2016
	180MG	A204097 003	Aug 19, 2016
RISING	60MG	A077081 006	Jul 21, 2011
	180MG	A077081 008	Jul 21, 2011
SCIEGEN PHARMS INC	60MG	A204507 002	Sep 16, 2015
	180MG	A204507 003	Sep 16, 2015
SUN PHARM INDS	180MG	A091567 006	Feb 06, 2012
TEVA	60MG	A076447 006	Apr 13, 2011
	180MG	A076447 008	Apr 13, 2011
UNIQUE	180MG	A210137 001	Aug 13, 2018
WOCKHARDT	60MG	A079112 004	Feb 08, 2012
	180MG	A079112 006	Feb 08, 2012

OTC DRUG PRODUCT LIST

FEXOFENADINE HYDROCHLORIDE

TABLET;ORAL

FEXOFENADINE HYDROCHLORIDE HIVES				
DR REDDYS LABS LTD	60MG	A076502	007	Apr 12, 2011
	180MG	A076502	009	Apr 12, 2011
RISING	60MG	A077081	007	Jul 21, 2011
	180MG	A077081	009	Jul 21, 2011
SCIEGEN PHARMS INC	60MG	A204507	004	Sep 16, 2015
	180MG	A204507	005	Sep 16, 2015
SUN PHARM INDS	180MG	A091567	005	Feb 06, 2012
TEVA	60MG	A076447	007	Apr 13, 2011
WOCKHARDT	60MG	A079112	003	Feb 08, 2012
	180MG	A079112	005	Feb 08, 2012

FEXOFENADINE HYDROCHLORIDE; PSEUDOEPHEDRINE HYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL

ALLEGRA-D 12 HOUR ALLERGY AND CONGESTION				
+! CHATTEM SANOFI	60MG;120MG	N020786	002	Jan 24, 2011
ALLEGRA-D 24 HOUR ALLERGY AND CONGESTION				
+! CHATTEM SANOFI	180MG;240MG	N021704	002	Jan 24, 2011
FEXOFENADINE HYDROCHLORIDE AND PSEUDOEPHEDRINE HYDROCHLORIDE				
AUROBINDO PHARMA	60MG;120MG	A209116	001	Oct 30, 2017
DR REDDYS	60MG;120MG	A076667	001	Nov 18, 2014
	60MG;120MG	A215434	001	May 31, 2022
DR REDDYS LABS LTD	180MG;240MG	A079043	002	Jun 22, 2011
SUN PHARM	60MG;120MG	A090818	001	Jan 29, 2015

FLUTICASONE FUROATE

SPRAY, METERED;NASAL

FLONASE SENSIMIST ALLERGY RELIEF				
+! HALEON US HOLDINGS	0.0275MG/SPRAY	N022051	002	Aug 02, 2016

FLUTICASONE PROPIONATE

SPRAY, METERED;NASAL

FLONASE ALLERGY RELIEF				
+ HALEON US HOLDINGS	0.05MG/SPRAY	N205434	001	Jul 23, 2014
FLUTICASONE PROPIONATE				
APOTEX	0.05MG/SPRAY	A208150	001	Feb 29, 2016
AUROBINDO PHARMA LTD	0.05MG/SPRAY	A217088	001	May 10, 2024
! HIKMA	0.05MG/SPRAY	A207957	001	May 26, 2016

GUAIFENESIN

TABLET, EXTENDED RELEASE;ORAL

GUAIFENESIN

ACTAVIS LABS FL	1.2GM	A091009	002	Sep 03, 2015
AMNEAL PHARMS	600MG	A207342	001	Jul 11, 2018
	1.2GM	A207342	002	Jul 11, 2018
AUROBINDO PHARMA	600MG	A210453	001	Oct 21, 2019
	1.2GM	A210453	002	Oct 21, 2019
DR REDDYS	600MG	A215932	001	Mar 15, 2022
	1.2GM	A215932	002	Mar 15, 2022
GRANULES	600MG	A213420	001	May 08, 2020
	1.2GM	A213420	002	May 08, 2020
GUARDIAN DRUG	600MG	A209215	001	Sep 06, 2017
	1.2GM	A209215	002	Sep 06, 2017
MARKSANS PHARMA	600MG	A217780	001	Aug 21, 2023
	1.2GM	A217780	002	Aug 21, 2023
OHM LABS INC	600MG	A209254	001	Jul 16, 2018
	1.2GM	A209254	002	Jul 16, 2018
PERRIGO R AND D	600MG	A078912	001	Nov 23, 2011
	1.2GM	A078912	002	Nov 05, 2020
MUCINEX				
+ RB HLTH	600MG	N021282	001	Jul 12, 2002
+!	1.2GM	N021282	002	Dec 18, 2002

GUAIFENESIN; PSEUDOEPHEDRINE HYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL

GUAIFENESIN AND PSEUDOEPHEDRINE HYDROCHLORIDE				
ACTAVIS LABS FL	600MG;60MG	A091071	001	May 27, 2015
	1.2GM;120MG	A091071	002	May 27, 2015
AUROBINDO PHARMA LTD	600MG;60MG	A213203	001	Mar 25, 2020
	1.2GM;120MG	A213203	002	Mar 25, 2020
DR REDDYS	600MG;60MG	A208369	001	Dec 29, 2017

OTC DRUG PRODUCT LIST

GUAIFENESIN; PSEUDOEPHEDRINE HYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL

GUAIFENESIN AND PSEUDOEPHEDRINE HYDROCHLORIDE

	1.2GM;120MG	A208369	002	Dec 29, 2017
L PERRIGO CO	600MG;60MG	A214407	001	Feb 01, 2022
	1.2GM;120MG	A214407	002	Feb 01, 2022
SUN PHARM INDS INC	600MG;60MG	A212542	001	Apr 28, 2020
	1.2GM;120MG	A212542	002	Apr 28, 2020
MUCINEX D				
+ RB HLTH	600MG;60MG	N021585	001	Jun 22, 2004
+!	1.2GM;120MG	N021585	002	Jun 22, 2004

IBUPROFEN

CAPSULE;ORAL

ADVIL LIQUI-GELS

+! HALEON US HOLDINGS	EQ 200MG FREE ACID AND POTASSIUM SALT	N020402	001	Apr 20, 1995
ADVIL MIGRAINE LIQUI-GELS				
+! HALEON US HOLDINGS	EQ 200MG FREE ACID AND POTASSIUM SALT	N020402	002	Mar 16, 2000

IBUPROFEN

AMNEAL PHARMS	EQ 200MG FREE ACID AND POTASSIUM SALT	A202300	001	Dec 23, 2011
ASCENT PHARMS INC	EQ 200MG FREE ACID AND POTASSIUM SALT	A206999	001	Dec 21, 2017
AUROBINDO PHARMA LTD	EQ 200MG FREE ACID AND POTASSIUM SALT	A207753	001	Jun 29, 2018
	EQ 200MG FREE ACID AND POTASSIUM SALT	A215777	001	Sep 14, 2023
BIONPHARMA	EQ 200MG FREE ACID AND POTASSIUM SALT	A078682	001	Mar 24, 2009
HUMANWELL PURACAP	EQ 200MG FREE ACID AND POTASSIUM SALT	A206568	001	Jun 21, 2016
MARKSANS PHARMA	EQ 200MG FREE ACID AND POTASSIUM SALT	A079205	001	Jun 26, 2009
P AND L DEV LLC	EQ 200MG FREE ACID AND POTASSIUM SALT	A077338	001	Jul 10, 2009
PATHEON SOFTGELS	EQ 200MG FREE ACID AND POTASSIUM SALT	A217236	001	Feb 13, 2024
SOFGEN PHARMS	EQ 200MG FREE ACID AND POTASSIUM SALT	A203599	001	Sep 07, 2016
STRIDES SOFTGELS	EQ 200MG FREE ACID AND POTASSIUM SALT	A204469	001	Mar 28, 2018

MIDOL LIQUID GELS

+! BIONPHARMA	200MG	N021472	001	Oct 18, 2002
---------------	-------	---------	-----	--------------

SUSPENSION;ORAL

CHILDREN'S ADVIL

HALEON US HOLDINGS	100MG/5ML	N020589	001	Jun 27, 1996
CHILDREN'S ADVIL-FLAVORED				
HALEON US HOLDINGS	100MG/5ML	N020589	002	Nov 07, 1997
CHILDREN'S IBUPROFEN				
PERRIGO	100MG/5ML	A074937	001	Dec 22, 1998

CHILDREN'S MOTRIN

+! KENVUE BRANDS	100MG/5ML	N020516	001	Jun 16, 1995
------------------	-----------	---------	-----	--------------

IBUPROFEN

ANNORA PHARMA	100MG/5ML	A210602	001	Nov 23, 2018
AUROBINDO PHARMA LTD	100MG/5ML	A209179	001	Apr 17, 2018
GUARDIAN DRUG	100MG/5ML	A210149	001	Aug 17, 2018
P AND L	100MG/5ML	A074916	001	Apr 30, 1999
PAI HOLDINGS PHARM	100MG/5ML	A214789	001	Nov 08, 2024
TARO	100MG/5ML	A209207	001	Jun 27, 2017

SUSPENSION/DROPS;ORAL

CHILDREN'S MOTRIN

+! KENVUE BRANDS	40MG/ML	N020603	001	Jun 10, 1996
------------------	---------	---------	-----	--------------

IBUPROFEN

GUARDIAN DRUG	40MG/ML	A210755	001	Sep 26, 2018
L PERRIGO CO	40MG/ML	A075217	001	Dec 16, 1998
TARO	40MG/ML	A217261	001	Aug 08, 2023
TRIS PHARMA INC	40MG/ML	A079058	001	Aug 31, 2009

INFANT'S ADVIL

+! HALEON US HOLDINGS	50MG/1.25ML	N020812	002	Jan 12, 2000
-----------------------	-------------	---------	-----	--------------

TABLET;ORAL

ADVIL

+! HALEON US HOLDINGS	200MG	N018989	001	May 18, 1984
-----------------------	-------	---------	-----	--------------

IBUPROFEN

ADAPTIS	200MG	A072299	001	Jul 01, 1988
AMNEAL PHARMS	200MG	A079233	001	Mar 18, 2014
AMNEAL PHARMS NY	200MG	A071333	001	Feb 17, 1987
	200MG	A072199	001	May 23, 1988
AUROBINDO PHARMA	200MG	A208865	001	Nov 08, 2017
AVEMA PHARMA	200MG	A076460	001	Nov 26, 2003
DR REDDYS	200MG	A075661	001	Dec 12, 2001
DR REDDYS LABS INC	200MG	A076117	001	Nov 20, 2001
ENDO OPERATIONS	200MG	A071229	001	Apr 01, 1987

OTC DRUG PRODUCT LIST

IBUPROFEN

TABLET;ORAL

IBUPROFEN					
GRANULES	200MG	A202312	001	Oct 07,	2016
GRANULES INDIA	200MG	A079174	001	Dec 10,	2010
LNK	200MG	A075010	001	Mar 01,	1999
	200MG	A075139	001	Mar 01,	1999
MARKSANS PHARMA	200MG	A091237	001	Feb 08,	2011
	200MG	A091239	001	Feb 01,	2011
MCNEIL	200MG	A073019	001	Mar 30,	1994
PERRIGO	200MG	A072096	001	Dec 08,	1987
! PERRIGO R AND D	200MG	A077349	001	Jun 21,	2005
SHANDONG XINHUA	200MG	A207095	001	May 05,	2017
	200MG	A207095	002	Aug 21,	2018
STRIDES PHARMA	200MG	A070481	001	Sep 24,	1986
	200MG	A079129	001	Mar 28,	2011
	200MG	A091355	001	Apr 04,	2011
	200MG	A206989	001	Jun 29,	2018
	200MG	A207052	001	May 30,	2017
JUNIOR STRENGTH ADVIL					
HALEON US HOLDINGS	100MG	N020267	002	Dec 13,	1996
JUNIOR STRENGTH MOTRIN					
KENVUE BRANDS	100MG	N020602	001	Jun 10,	1996
MOTRIN IB					
+ KENVUE BRANDS	200MG	N019012	003	Dec 17,	1990
TABLET, CHEWABLE;ORAL					
CHILDREN'S ADVIL					
HALEON US HOLDINGS	50MG	N020944	001	Dec 18,	1998
IBUPROFEN					
! PERRIGO	100MG	A076359	002	Jan 16,	2004
JUNIOR STRENGTH ADVIL					
HALEON US HOLDINGS	100MG	N020944	002	Dec 18,	1998

IBUPROFEN SODIUM

TABLET;ORAL

ADVIL

+! HALEON US HOLDINGS	EQ 200MG BASE	N201803	001	Jun 12,	2012
-----------------------	---------------	---------	-----	---------	------

IBUPROFEN; PHENYLEPHRINE HYDROCHLORIDE

TABLET;ORAL

ADVIL CONGESTION RELIEF

+! HALEON US HOLDINGS	200MG;10MG	N022565	001	May 27,	2010
-----------------------	------------	---------	-----	---------	------

IBUPROFEN AND PHENYLEPHRINE HYDROCHLORIDE

PERRIGO R AND D	200MG;10MG	A203200	001	Jul 03,	2014
-----------------	------------	---------	-----	---------	------

IBUPROFEN; PSEUDOEPHEDRINE HYDROCHLORIDE

CAPSULE;ORAL

ADVIL COLD AND SINUS

+! HALEON US HOLDINGS	EQ 200MG FREE ACID AND POTASSIUM SALT;30MG	N021374	001	May 30,	2002
-----------------------	--	---------	-----	---------	------

IBUPROFEN AND PSEUDOEPHEDRINE HYDROCHLORIDE

AUROBINDO PHARMA	EQ 200MG FREE ACID AND POTASSIUM SALT;30MG	A209235	001	Dec 01,	2017
------------------	--	---------	-----	---------	------

SUSPENSION;ORAL

CHILDREN'S ADVIL COLD

HALEON US HOLDINGS	100MG/5ML;15MG/5ML	N021373	001	Apr 18,	2002
--------------------	--------------------	---------	-----	---------	------

CHILDREN'S MOTRIN COLD

+! KENVUE BRANDS	100MG/5ML;15MG/5ML	N021128	001	Aug 01,	2000
------------------	--------------------	---------	-----	---------	------

TABLET;ORAL

ADVIL COLD AND SINUS

+! HALEON US HOLDINGS	200MG;30MG	N019771	001	Sep 19,	1989
-----------------------	------------	---------	-----	---------	------

IBUPROFEN AND PSEUDOEPHEDRINE HYDROCHLORIDE

DR REDDYS LABS LTD	200MG;30MG	A077628	001	Aug 14,	2006
--------------------	------------	---------	-----	---------	------

IBUPROHM COLD AND SINUS

OHM LABS	200MG;30MG	A074567	001	Apr 17,	2001
----------	------------	---------	-----	---------	------

SINE-AID IB

KENVUE BRANDS	200MG;30MG	N019899	001	Dec 31,	1992
---------------	------------	---------	-----	---------	------

IODINE POVACRYLEX; ISOPROPYL ALCOHOL

SPONGE;TOPICAL

DURAPREP

+! 3M	EQ 0.7% IODINE;74% (6ML)	N021586	001	Sep 29,	2006
-------	--------------------------	---------	-----	---------	------

+!	EQ 0.7% IODINE;74% (26ML)	N021586	002	Sep 29,	2006
----	---------------------------	---------	-----	---------	------

OTC DRUG PRODUCT LIST

ISOPROPYL ALCOHOL

SOLUTION;TOPICAL

ZURAGARD

+! ZUREX PHARMA 70% N210872 001 Apr 26, 2019

IVERMECTIN

LOTION;TOPICAL

IVERMECTIN

TARO 0.5% A210720 001 May 06, 2020

SKLICE

+! ARBOR PHARMS LLC 0.5% N202736 001 Feb 07, 2012

KETOCONAZOLE

SHAMPOO;TOPICAL

NIZORAL ANTI-DANDRUFF

+! KRAMER 1% N020310 001 Oct 10, 1997

KETOTIFEN FUMARATE

SOLUTION/DROPS;OPHTHALMIC

ALAWAY

+! BAUSCH AND LOMB EQ 0.025% BASE N021996 001 Dec 01, 2006

CHILDREN'S ALAWAY

+ BAUSCH AND LOMB EQ 0.025% BASE N021996 002 Feb 11, 2015

KETOTIFEN FUMARATE

APOTEX INC EQ 0.025% BASE A077354 001 May 09, 2006

SENTISS EQ 0.025% BASE A077958 001 Jul 26, 2007

UNICHEM EQ 0.025% BASE A204059 001 Jun 01, 2020

ZADITOR

! ALCON PHARMS LTD EQ 0.025% BASE A077200 001 Sep 02, 2008

LANSOPRAZOLE

CAPSULE, DELAYED REL PELLETS;ORAL

LANSOPRAZOLE

DR REDDYS LABS LTD 15MG A202194 001 May 18, 2012

GLENMARK PHARMS INC 15MG A202727 001 May 18, 2012

NATCO 15MG A203306 001 Jan 13, 2016

PERRIGO R AND D 15MG A202319 001 May 18, 2012

PREVACID 24 HR

+! PERRIGO PHARMA INTL 15MG N022327 001 May 18, 2009

TABLET, ORALLY DISINTEGRATING, DELAYED RELEASE;ORAL

LANSOPRAZOLE

DEXCEL 15MG N208025 001 Jun 07, 2016

LEVOCETIRIZINE DIHYDROCHLORIDE

SOLUTION;ORAL

XYZAL ALLERGY 24HR

+! CHATTEM SANOFI 2.5MG/5ML N209090 001 Jan 31, 2017

TABLET;ORAL

LEVOCETIRIZINE DIHYDROCHLORIDE

APOTEX 5MG A211443 001 Apr 21, 2021

DR REDDYS 5MG A210375 001 Jan 19, 2018

HETERO LABS LTD III 5MG A213513 001 Apr 29, 2020

MICRO LABS 5MG A211551 001 Nov 20, 2018

PERRIGO R AND D 5MG A211983 001 Mar 28, 2019

XYZAL ALLERGY 24HR

+! CHATTEM SANOFI 5MG N209089 001 Jan 31, 2017

LEVONORGESTREL

TABLET;ORAL

ATHENTIA NEXT

AUROBINDO PHARMA 1.5MG A206867 001 Dec 08, 2015

FALLBACK SOLO

LUPIN LTD 1.5MG A201446 001 Jun 19, 2014

HER STYLE

NOVAST LABS 1.5MG A207976 001 Mar 11, 2016

LEVONORGESTREL

GLENMARK PHARMS LTD 1.5MG A207044 001 Mar 25, 2016

LABORATOIRE HRA 1.5MG A204044 001 Jul 03, 2018

NAARI PTE LTD 1.5MG A202380 001 May 29, 2015

1.5MG A207660 001 May 02, 2019

NOVEL LABS INC 1.5MG A202508 001 Feb 22, 2013

PERRIGO R AND D 1.5MG A202334 001 Aug 20, 2014

1.5MG A202739 001 Oct 31, 2014

1.5MG A205329 001 Sep 18, 2018

OTC DRUG PRODUCT LIST

LEVONORGESTREL

TABLET; ORAL

OPCICON ONE-STEP

SUN PHARM

1.5MG

A202635 001 Sep 11, 2014

PLAN B ONE-STEP

+! FDN CONSUMER

1.5MG

N021998 001 Jul 10, 2009

LOPERAMIDE HYDROCHLORIDE

CAPSULE; ORAL

LOPERAMIDE HYDROCHLORIDE

+ BIONPHARMA

1MG

N021855 001 Aug 04, 2005

+!

2MG

N021855 002 Aug 04, 2005

STRIDES SOFTGELS

2MG

A213070 001 Aug 11, 2021

SOLUTION; ORAL

IMODIUM A-D

+! KENVUE BRANDS

1MG/5ML

N019487 001 Mar 01, 1988

LOPERAMIDE HYDROCHLORIDE

WOCKHARDT BIO AG

1MG/5ML

A074730 001 Aug 28, 1997

SUSPENSION; ORAL

IMODIUM A-D

+! KENVUE BRANDS

1MG/7.5ML

N019487 002 Jul 08, 2004

LOPERAMIDE HYDROCHLORIDE

PERRIGO R AND D

1MG/7.5ML

A091292 001 May 20, 2011

TABLET; ORAL

IMODIUM A-D

+! KENVUE BRANDS

2MG

N019860 001 Nov 22, 1989

LOPERAMIDE HYDROCHLORIDE

AUROBINDO PHARMA

2MG

A206548 001 Dec 15, 2015

L PERRIGO CO

2MG

A075232 001 Jan 06, 2000

LNK

2MG

A076497 001 Jun 10, 2003

OHM LABS

2MG

A074091 001 Dec 10, 1992

LOPERAMIDE HYDROCHLORIDE; SIMETHICONE

TABLET; ORAL

IMODIUM MULTI-SYMPOM RELIEF

+! KENVUE BRANDS

2MG;125MG

N021140 001 Nov 30, 2000

LOPERAMIDE HYDROCHLORIDE AND SIMETHICONE

AUROBINDO PHARMA

2MG;125MG

A211059 001 Dec 14, 2020

LTD

BIONPHARMA

2MG;125MG

A213484 001 Sep 10, 2021

GUARDIAN DRUG

2MG;125MG

A214541 001 May 27, 2021

HETERO LABS LTD V

2MG;125MG

A211438 001 Jun 17, 2021

PERRIGO R AND D

2MG;125MG

A209837 001 Sep 05, 2018

LORATADINE

CAPSULE; ORAL

CLARITIN

+! BAYER HEALTHCARE

10MG

N021952 001 Jun 16, 2008

LLC

LORATADINE

BIONPHARMA

10MG

A202538 001 Dec 21, 2018

MARKSANS PHARMA

10MG

A206214 001 Sep 23, 2016

SUSPENSION; ORAL

LORATADINE

+! TARO

1MG/ML

N021734 001 Oct 04, 2005

SYRUP; ORAL

CLARITIN

+! BAYER HEALTHCARE

1MG/ML

N020641 002 Nov 27, 2002

LLC

LORATADINE

AUROBINDO PHARMA

1MG/ML

A208931 001 Jun 29, 2018

LTD

HETERO LABS LTD III

1MG/ML

A210409 001 May 07, 2021

LANNETT CO INC

1MG/ML

A077421 001 Jun 29, 2006

PERRIGO

1MG/ML

A075728 001 Aug 20, 2004

TARO

1MG/ML

A076805 001 Aug 20, 2004

1MG/ML

A201865 001 Jul 31, 2015

WOCKHARDT BIO AG

1MG/ML

A075815 001 Aug 20, 2004

TABLET; ORAL

CLARITIN

+! BAYER HEALTHCARE

10MG

N019658 002 Nov 27, 2002

LLC

CLARITIN HIVES RELIEF

+! BAYER HEALTHCARE

10MG

N019658 003 Nov 19, 2003

LLC

OTC DRUG PRODUCT LIST

LORATADINE

TABLET; ORAL

LORATADINE

ANDA REPOSITORY	10MG	A207569	001	Mar 12, 2019
APOTEX INC	10MG	A076471	001	Feb 14, 2006
AUROBINDO PHARMA LTD	10MG	A208314	001	Apr 16, 2018
GRANULES	10MG	A210722	001	Dec 23, 2019
HETERO LABS LTD V	10MG	A211718	001	Jul 28, 2023
MARKSANS PHARMA	10MG	A219223	001	Nov 21, 2024
MYLAN	10MG	A076154	001	Aug 20, 2003
PERRIGO	10MG	A076301	001	Jun 25, 2004
PLD ACQUISITIONS LLC	10MG	A075209	001	Jan 21, 2003
SUN PHARM INDS LTD	10MG	A076134	001	Aug 18, 2003
UNIQUE PHARM	10MG	A214684	001	Jan 07, 2021

TABLET, CHEWABLE; ORAL

CHILDREN'S CLARITIN

+ BAYER HEALTHCARE LLC	5MG	N021891	001	Aug 23, 2006
------------------------	-----	---------	-----	--------------

CLARITIN

+! BAYER HEALTHCARE LLC	10MG	N021891	002	Nov 21, 2018
-------------------------	------	---------	-----	--------------

LORATADINE

PERRIGO PHARMA INTL	5MG	A210033	001	Jun 12, 2019
SUN PHARM	5MG	A210088	001	Apr 16, 2018

TABLET, ORALLY DISINTEGRATING; ORAL

ALAVERT

FDN CONSUMER	10MG	N021375	001	Dec 19, 2002
--------------	------	---------	-----	--------------

CLARITIN HIVES RELIEF REDITAB

+! BAYER HEALTHCARE LLC	10MG	N020704	003	Nov 19, 2003
-------------------------	------	---------	-----	--------------

CLARITIN REDITABS

+! BAYER HEALTHCARE LLC	5MG	N021993	001	Dec 12, 2006
-------------------------	-----	---------	-----	--------------

+!	10MG	N020704	002	Nov 27, 2002
----	------	---------	-----	--------------

LORATADINE

AUROBINDO PHARMA LTD	10MG	A208477	001	Apr 11, 2018
PERRIGO PHARMA INTL	10MG	A076011	001	Sep 29, 2003
RUBICON	10MG	A214280	001	Sep 10, 2020
TENSHI	5MG	A212795	001	Sep 18, 2020
	10MG	A213294	001	Oct 30, 2020

LORATADINE REDIDOSE

SUN PHARM INDS LTD	10MG	A077153	001	Apr 11, 2007
--------------------	------	---------	-----	--------------

LORATADINE; PSEUDOEPHEDRINE SULFATE

TABLET, EXTENDED RELEASE; ORAL

CLARITIN-D

+! BAYER HEALTHCARE LLC	5MG;120MG	N019670	002	Nov 27, 2002
-------------------------	-----------	---------	-----	--------------

CLARITIN-D 24 HOUR

+! BAYER HEALTHCARE LLC	10MG;240MG	N020470	002	Nov 27, 2002
-------------------------	------------	---------	-----	--------------

LORATADINE AND PSEUDOEPHEDRINE SULFATE

BIONPHARMA	10MG;240MG	A218017	001	May 14, 2024
P AND L	10MG;240MG	A075706	001	Feb 21, 2003
PERRIGO PHARMA INTL	5MG;120MG	A076050	001	Jan 30, 2003
	10MG;240MG	A075989	001	Mar 04, 2004
SUN PHARM INDS LTD	10MG;240MG	A076557	001	Sep 22, 2004

MENTHOL; METHYL SALICYLATE

PATCH; TOPICAL

SALONPAS

+! HISAMITSU PHARM CO	3%;10%	N022029	001	Feb 20, 2008
+	3%;10%	N022029	002	Nov 05, 2012

MICONAZOLE NITRATE

CREAM; TOPICAL, VAGINAL

MICONAZOLE 3 COMBINATION PACK

PERRIGO	2%,4%	A076357	001	Mar 30, 2004
---------	-------	---------	-----	--------------

MONISTAT 3 COMBINATION PACK

+ MEDTECH PRODUCTS	2%,4%	N021261	003	Jun 17, 2003
--------------------	-------	---------	-----	--------------

MONISTAT 3 COMBINATION PACK (PREFILLED)

+! MEDTECH PRODUCTS	2%,4%	N021261	001	Feb 02, 2001
---------------------	-------	---------	-----	--------------

OTC DRUG PRODUCT LIST

MICONAZOLE NITRATE

CREAM;VAGINAL

MICONAZOLE 3

TARO 4%

A076773 001 Mar 02, 2005

MICONAZOLE 7

P AND L 2%

A074164 001 Mar 29, 1996

MICONAZOLE NITRATE

PERRIGO 2%

A074760 001 May 15, 1997

PERRIGO R AND D 4%

A091366 001 Jan 15, 2010

TARO 2%

A074444 001 Jan 13, 1997

MONISTAT 3

+! MEDTECH PRODUCTS 4%

N020827 001 Mar 30, 1998

MONISTAT 7

+! MEDTECH PRODUCTS 2%

N017450 002 Feb 15, 1991

CREAM, INSERT;TOPICAL, VAGINAL

MICONAZOLE NITRATE

PERRIGO R AND D 2%,1.2GM

A079114 001 Jun 02, 2010

MONISTAT 1 COMBINATION PACK

+! MEDTECH PRODUCTS 2%,1.2GM

N021308 001 Jun 29, 2001

CREAM, SUPPOSITORY;TOPICAL, VAGINAL

M-ZOLE 3 COMBINATION PACK

COSETTE 2%,200MG

A074926 001 Apr 16, 1999

MICONAZOLE NITRATE COMBINATION PACK

L PERRIGO CO 2%,200MG

A075329 001 Apr 20, 1999

MONISTAT 3 COMBINATION PACK

+! MEDTECH PRODUCTS 2%,200MG

N020670 002 Apr 16, 1996

MONISTAT 7 COMBINATION PACK

+! MEDTECH PRODUCTS 2%,100MG

N020288 002 Apr 26, 1993

SUPPOSITORY;VAGINAL

MICONAZOLE NITRATE

COSETTE 100MG

A073507 001 Nov 19, 1993

MONISTAT 7

+! MEDTECH PRODUCTS 100MG

N018520 002 Feb 15, 1991

MINOXIDIL

AEROSOL, FOAM;TOPICAL

MEN'S ROGAINE

+! KENVUE BRANDS 5%

N021812 001 Jan 20, 2006

MINOXIDIL

PERRIGO PHARMA INTL 5%

A091344 001 Apr 28, 2011

MINOXIDIL (FOR MEN)

AUROBINDO PHARMA 5%

A218616 001 Apr 22, 2024

P AND L 5%

A208092 001 Feb 17, 2017

TARO 5%

A209074 001 Dec 31, 2018

MINOXIDIL (FOR WOMEN)

AUROBINDO PHARMA 5%

A218616 002 Apr 22, 2024

P AND L 5%

A208092 002 Jul 27, 2017

TARO 5%

A209074 002 Apr 22, 2019

WOMEN'S ROGAINE

+! KENVUE BRANDS 5%

N021812 002 Feb 28, 2014

SOLUTION;TOPICAL

MINOXIDIL (FOR MEN)

L PERRIGO CO 2%

A075357 001 Jul 30, 1999

P AND L 2%

A074588 001 Apr 05, 1996

MINOXIDIL (FOR WOMEN)

L PERRIGO CO 2%

A075357 002 Jul 30, 1999

TARO 2%

A218175 001 Nov 26, 2024

MINOXIDIL EXTRA STRENGTH (FOR MEN)

AUROBINDO PHARMA 5%

A075438 001 Feb 27, 2003

LTD

P AND L 5%

A075518 001 Nov 17, 2000

PERRIGO 5%

A075598 001 Jun 13, 2001

ROGAINE (FOR MEN)

+! KENVUE BRANDS 2%

N019501 002 Feb 09, 1996

ROGAINE (FOR WOMEN)

+! KENVUE BRANDS 2%

N019501 003 Feb 09, 1996

ROGAINE EXTRA STRENGTH (FOR MEN)

+! KENVUE BRANDS 5%

N020834 001 Nov 14, 1997

THEROXIDIL

PURE SOURCE 2%

A078176 001 Nov 09, 2007

5%

A076239 001 Aug 24, 2004

OTC DRUG PRODUCT LIST

MOMETASONE FUROATE

SPRAY, METERED;NASAL

MOMETASONE FUROATE

AUROBINDO PHARMA 0.05MG/SPRAY

A217498 001 Nov 22, 2024

NASONEX 24HR ALLERGY

+! PERRIGO PHARMA INTL 0.05MG/SPRAY

N215712 001 Mar 17, 2022

NALOXONE HYDROCHLORIDE

SPRAY, METERED;NASAL

NALOXONE HYDROCHLORIDE

AMNEAL 4MG/SPRAY

A217992 001 Apr 23, 2024

PADAGIS ISRAEL 4MG/SPRAY

A211951 001 Jun 21, 2022

TEVA PHARMS USA 4MG/SPRAY

A209522 001 Apr 19, 2019

NARCAN

+! EMERGENT 4MG/SPRAY

N208411 001 Nov 18, 2015

RIVIVE

+! HARM REDUCTION 3MG/SPRAY

N217722 001 Jul 28, 2023

THERP

NAPHAZOLINE HYDROCHLORIDE; PHENIRAMINE MALEATE

SOLUTION/DROPS;OPHTHALMIC

NAPHAZOLINE HYDROCHLORIDE AND PHENIRAMINE MALEATE

ALTAIRE PHARMS INC 0.02675%;0.315%

A078208 001 Sep 27, 2010

RISING 0.025%;0.3%

A202795 001 Jan 24, 2013

NAPHCN-A

+! ALCON 0.025%;0.3%

N020226 001 Jun 08, 1994

OPCON-A

+! BAUSCH AND LOMB 0.02675%;0.315%

N020065 001 Jun 08, 1994

VISINE

+! KENVUE BRANDS 0.025%;0.3%

N020485 001 Jan 31, 1996

NAPROXEN SODIUM

CAPSULE;ORAL

NAPROXEN SODIUM

+! BIONPHARMA EQ 200MG BASE

N021920 001 Feb 17, 2006

CATALENT EQ 200MG BASE

A202807 001 Jan 04, 2019

PATHEON SOFTGELS EQ 200MG BASE

A214463 001 Jan 10, 2023

PURACAP PHARM LLC EQ 200MG BASE

A208363 001 Mar 15, 2018

TABLET;ORAL

ALEVE

+! BAYER 220MG

N020204 002 Jan 11, 1994

NAPROXEN SODIUM

AMNEAL PHARMS NY 220MG

A079096 001 Dec 16, 2008

AUROBINDO PHARMA 220MG

A205497 001 Mar 18, 2016

LTD

CONTRACT PHARMACAL 220MG

A074635 001 Jan 13, 1997

DR REDDYS LABS INC 220MG

A075168 001 Jul 28, 1998

GRANULES INDIA 220MG

A091353 001 Sep 20, 2011

HETERO LABS LTD V 220MG

A211065 001 Oct 28, 2022

LNK INTL INC 220MG

A204872 001 Jan 23, 2017

MARKSANS PHARMA 220MG

A090545 001 Mar 16, 2011

NOVELGENIX THERAPS 220MG

A207612 001 Nov 16, 2018

PERRIGO 220MG

A074661 001 Jan 13, 1997

YICHANG HUMANWELL 220MG

A212033 001 Aug 30, 2019

NAPROXEN SODIUM; PSEUDOEPHEDRINE HYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL

NAPROXEN SODIUM AND PSEUDOEPHEDRINE HYDROCHLORIDE

AUROBINDO PHARMA 220MG;120MG

A211360 001 Jun 01, 2022

DR REDDYS LABS INC 220MG;120MG

A077381 001 Sep 27, 2006

! PERRIGO 220MG;120MG

A076518 001 Mar 17, 2004

NICOTINE

FILM, EXTENDED RELEASE;TRANSDERMAL

HABITROL

+ DR REDDYS LABS SA 7MG/24HR

N020076 004 Nov 12, 1999

+ 14MG/24HR

N020076 005 Nov 12, 1999

+! 21MG/24HR

N020076 006 Nov 12, 1999

NICODERM CQ

+ CHATTEM SANOFI 7MG/24HR

N020165 006 Aug 02, 1996

+ 14MG/24HR

N020165 005 Aug 02, 1996

+! 21MG/24HR

N020165 004 Aug 02, 1996

NICOTINE

DIFGEN PHARMS 7MG/24HR

A074612 002 Jul 28, 2003

14MG/24HR

A074612 003 Oct 20, 1997

OTC DRUG PRODUCT LIST

NICOTINE

FILM, EXTENDED RELEASE;TRANSDERMAL

NICOTINE

21MG/24HR

A074612 001 Oct 20, 1997

NICOTINE POLACRILEX

GUM, CHEWING;BUCCAL

NICORETTE

+ HALEON US HOLDINGS

EQ 2MG BASE

N018612 002 Feb 09, 1996

+

EQ 2MG BASE

N018612 004 Sep 25, 2000

+!

EQ 4MG BASE

N020066 002 Feb 09, 1996

+

EQ 4MG BASE

N020066 004 Sep 25, 2000

NICORETTE (MINT)

+ HALEON US HOLDINGS

EQ 2MG BASE

N018612 003 Dec 23, 1998

+

EQ 4MG BASE

N020066 003 Dec 23, 1998

NICOTINE POLACRILEX

L PERRIGO CO

EQ 2MG BASE

A076775 001 Sep 16, 2004

EQ 2MG BASE

A076777 001 Sep 16, 2004

EQ 4MG BASE

A076779 001 Sep 16, 2004

EQ 4MG BASE

A076789 001 Sep 16, 2004

P AND L

EQ 2MG BASE

A074507 001 Mar 15, 1999

EQ 2MG BASE

A076569 001 Jul 29, 2004

EQ 2MG BASE

A078699 001 Dec 29, 2008

EQ 2MG BASE

A079044 001 Jul 08, 2009

EQ 2MG BASE

A079216 001 Jul 08, 2009

EQ 2MG BASE

A204794 001 May 10, 2016

EQ 4MG BASE

A074707 001 Mar 19, 1999

EQ 4MG BASE

A076568 002 Jul 29, 2004

EQ 4MG BASE

A078697 001 Dec 29, 2008

EQ 4MG BASE

A079038 001 Jul 08, 2009

EQ 4MG BASE

A079219 001 Jul 08, 2009

EQ 4MG BASE

A204833 001 Feb 26, 2016

PERRIGO R AND D

EQ 2MG BASE

A078325 001 Oct 30, 2006

EQ 2MG BASE

A078547 001 May 24, 2007

EQ 2MG BASE

A091349 001 Jul 20, 2011

EQ 2MG BASE

A206394 001 Dec 15, 2016

EQ 4MG BASE

A078326 001 Oct 30, 2006

EQ 4MG BASE

A078546 001 May 24, 2007

EQ 4MG BASE

A091354 001 Jul 20, 2011

EQ 4MG BASE

A206393 001 Dec 15, 2016

TROCHE/LOZENGE;ORAL

NICORETTE

+ HALEON US HOLDINGS

EQ 2MG BASE

N021330 001 Oct 31, 2002

+

EQ 2MG BASE

N022360 001 May 18, 2009

+!

EQ 4MG BASE

N021330 002 Oct 31, 2002

+!

EQ 4MG BASE

N022360 002 May 18, 2009

NICOTINE POLACRILEX

AUROBINDO PHARMA

EQ 2MG BASE

A213266 001 Aug 03, 2021

EQ 4MG BASE

A213266 002 Aug 03, 2021

AUROBINDO PHARMA

EQ 2MG BASE

A215357 001 May 13, 2022

LTD

EQ 4MG BASE

A215357 002 May 13, 2022

DR REDDYS LABS SA

EQ 2MG BASE

A212796 001 Jan 08, 2020

EQ 2MG BASE

A212983 001 Feb 21, 2020

EQ 2MG BASE

A213233 001 Aug 04, 2020

EQ 4MG BASE

A212796 002 Jan 08, 2020

EQ 4MG BASE

A212983 002 Feb 21, 2020

EQ 4MG BASE

A213233 002 Aug 04, 2020

P AND L

EQ 2MG BASE

A208875 001 Oct 31, 2019

EQ 2MG BASE

A209206 001 Jun 26, 2018

EQ 2MG BASE

A209519 001 Jul 02, 2018

EQ 2MG BASE

A209520 001 Oct 31, 2019

EQ 2MG BASE

A210711 001 Oct 31, 2019

EQ 2MG BASE

A210712 001 Sep 06, 2019

EQ 2MG BASE

A212056 001 Jul 26, 2019

EQ 2MG BASE

A212057 001 May 14, 2020

EQ 4MG BASE

A208875 002 Oct 31, 2019

EQ 4MG BASE

A209206 002 Jun 26, 2018

EQ 4MG BASE

A209519 002 Jul 02, 2018

EQ 4MG BASE

A209520 002 Oct 31, 2019

EQ 4MG BASE

A210711 002 Oct 31, 2019

EQ 4MG BASE

A210712 002 Sep 06, 2019

EQ 4MG BASE

A212056 002 Jul 26, 2019

OTC DRUG PRODUCT LIST

NICOTINE POLACRILEX

TROCHE/LOZENGE;ORAL

NICOTINE POLACRILEX

	EQ 4MG BASE	A212057	002	May 14, 2020
PERRIGO R AND D	EQ 2MG BASE	A077007	001	Jan 31, 2006
	EQ 2MG BASE	A090821	001	Jul 10, 2009
	EQ 4MG BASE	A077007	002	Jan 31, 2006
	EQ 4MG BASE	A090821	002	Jul 10, 2009
PLD ACQUISITIONS	EQ 2MG BASE	A207868	001	Feb 07, 2019
	EQ 4MG BASE	A207868	002	Feb 07, 2019
PPI-DAC	EQ 2MG BASE	A090711	001	Jul 10, 2009
	EQ 2MG BASE	A203690	001	Oct 09, 2012
	EQ 4MG BASE	A090711	002	Jul 10, 2009
	EQ 4MG BASE	A203690	002	Oct 09, 2012

NIZATIDINE

TABLET;ORAL

AXID AR

+!	HALEON US HOLDINGS	75MG	N020555	001	May 09, 1996
----	--------------------	------	---------	-----	--------------

NORGESTREL

TABLET;ORAL

OPILL

+!	LABORATOIRE HRA	0.075MG	N017031	001	
----	-----------------	---------	---------	-----	--

OLOPATADINE HYDROCHLORIDE

SOLUTION/DROPS;OPHTHALMIC

OLOPATADINE HYDROCHLORIDE

ALEMBIC	EQ 0.1% BASE	A209919	001	Dec 07, 2018	
	EQ 0.2% BASE	A209420	001	Apr 29, 2019	
APOTEX	EQ 0.1% BASE	A078350	001	Dec 07, 2015	
	EQ 0.2% BASE	A090918	001	Dec 05, 2017	
BARR LABS INC	EQ 0.2% BASE	A090848	001	Jul 13, 2015	
EUGIA PHARMA	EQ 0.1% BASE	A204812	001	Dec 18, 2015	
	EQ 0.2% BASE	A209995	001	Apr 04, 2019	
GLAND PHARMA LTD	EQ 0.1% BASE	A209619	001	Aug 02, 2019	
	EQ 0.2% BASE	A209752	001	May 20, 2020	
GLENMARK PHARMS INC	EQ 0.1% BASE	A200810	001	Jun 28, 2017	
SOMERSET	EQ 0.2% BASE	A215006	001	Dec 18, 2024	
SOMERSET THERAPS	EQ 0.1% BASE	A206306	001	Dec 07, 2015	
LLC					
USV	EQ 0.1% BASE	A203152	001	Dec 07, 2015	
PATADAY ONCE DAILY RELIEF					
+!	ALCON LABS INC	EQ 0.2% BASE	N021545	001	Dec 22, 2004
+!		EQ 0.7% BASE	N206276	001	Jan 30, 2015
PATADAY TWICE DAILY RELIEF					
+!	ALCON LABS INC	EQ 0.1% BASE	N020688	001	Dec 18, 1996

OMEPRAZOLE

TABLET, DELAYED RELEASE;ORAL

OMEPRAZOLE

APOTEX	20MG	A210070	001	Feb 11, 2019	
+!	DEXCEL PHARMA	20MG	N022032	001	Dec 04, 2007
	DR REDDYS	20MG	A207740	001	Nov 05, 2018
	SUN PHARM	20MG	A207891	001	Oct 12, 2018

TABLET, ORALLY DISINTEGRATING, DELAYED RELEASE;ORAL

OMEPRAZOLE

+!	DEXCEL	20MG	N209400	001	Jul 05, 2017
----	--------	------	---------	-----	--------------

OMEPRAZOLE MAGNESIUM

CAPSULE, DELAYED RELEASE;ORAL

OMEPRAZOLE MAGNESIUM

!	DR REDDYS LABS LTD	EQ 20MG BASE	A078878	001	Jun 05, 2009
	L PERRIGO CO	EQ 20MG BASE	A216096	001	May 24, 2022

TABLET, DELAYED RELEASE;ORAL

OMEPRAZOLE MAGNESIUM

AUROBINDO PHARMA	EQ 20MG BASE	A206877	001	Jun 06, 2018	
HETERO LABS LTD III	EQ 20MG BASE	A211732	001	Mar 25, 2020	
P AND L	EQ 20MG BASE	A206582	001	Jun 01, 2020	
PRILOSEC OTC					
+!	ASTRAZENECA	EQ 20MG BASE	N021229	001	Jun 20, 2003

OTC DRUG PRODUCT LIST

OMEPRAZOLE; SODIUM BICARBONATE

CAPSULE; ORAL

OMEPRAZOLE AND SODIUM BICARBONATE

AUROBINDO PHARMA 20MG; 1.1GM

A204923 001 Nov 07, 2016

PERRIGO R AND D 20MG; 1.1GM

A201361 001 Jul 15, 2016

ZYDUS 20MG; 1.1GM

A203345 001 Mar 16, 2018

ZEGERID OTC

+! RILEY CONSUMER 20MG; 1.1GM

N022281 001 Dec 01, 2009

FOR SUSPENSION; ORAL

OMEPRAZOLE AND SODIUM BICARBONATE

DR REDDYS 20MG/PACKET; 1.68GM/PACKET

A217784 001 May 09, 2024

ZEGERID OTC

+! RILEY CONSUMER 20MG/PACKET; 1.68GM/PACKET

N022283 001 Jun 17, 2013

ORLISTAT

CAPSULE; ORAL

ALLI

+! HALEON US HOLDINGS 60MG

N021887 001 Feb 07, 2007

OXYBUTYRNIN

FILM, EXTENDED RELEASE; TRANSDERMAL

OXYTROL FOR WOMEN

+! ABBVIE 3.9MG/24HR

N202211 001 Jan 25, 2013

OXYMETAZOLINE HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

VISINE L.R.

+! KENVUE BRANDS 0.025%

N019407 001 Mar 31, 1989

PERMETHRIN

LOTION; TOPICAL

NIX

+! MEDTECH PRODUCTS 1%

N019918 001 May 02, 1990

PERMETHRIN

ACTAVIS MID 1%

A075014 001 Mar 28, 2000

ATLANTIC

PERRIGO NEW YORK 1%

A076090 001 Dec 20, 2001

POLYETHYLENE GLYCOL 3350

FOR SOLUTION; ORAL

MIRALAX

+! BAYER HEALTHCARE 17GM/SCOOPFUL

N022015 001 Oct 06, 2006

LLC

POLYETHYLENE GLYCOL 3350

ANI PHARMS 17GM/SCOOPFUL

A202850 001 Dec 15, 2015

ANNORA PHARMA 17GM/SCOOPFUL

A214990 001 Apr 14, 2021

AUROBINDO PHARMA 17GM/SCOOPFUL

A209017 001 Apr 09, 2018

LTD

ELYSIUM 17GM/SCOOPFUL

A202071 001 Dec 28, 2012

LGM PHARMA 17GM/SCOOPFUL

A090812 001 Oct 07, 2009

MYLAN 17GM/PACKET

A078915 001 Oct 06, 2009

17GM/SCOOPFUL

A078915 002 Oct 06, 2009

NOVEL LABS INC 17GM/SCOOPFUL

A091077 001 Oct 06, 2009

NUVO PHARMS INC 17GM/SCOOPFUL

A206105 001 Oct 28, 2016

PPI-DAC 17GM/PACKET

A090685 001 Oct 06, 2009

17GM/SCOOPFUL

A090685 002 Oct 06, 2009

STRIDES PHARMA 17GM/SCOOPFUL

A079214 001 Jan 31, 2013

17GM/SCOOPFUL

A203928 001 Aug 24, 2016

17GM/PACKET

A203928 002 Aug 24, 2016

POTASSIUM IODIDE

SOLUTION; ORAL

POTASSIUM IODIDE

! MISSION PHARMACAL 65MG/ML

A206211 001 Mar 24, 2016

TABLET; ORAL

IOSAT

+ ANBEX 65MG

N018664 002 May 12, 2011

+! 130MG

N018664 001 Oct 14, 1982

THYROSAFE

! BTG INTL 65MG

A076350 001 Sep 10, 2002

OTC DRUG PRODUCT LIST

POVIDONE-IODINE

SOLUTION;TOPICAL

POVIDONE IODINE

+! ALLEGIANCE HLTHCARE 1%

N019522 001 Mar 31, 1989

PSEUDOEPHEDRINE HYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL

PSEUDOEPHEDRINE HYDROCHLORIDE

AUROBINDO PHARMA 120MG

A209008 001 Jun 09, 2017

! L PERRIGO CO 120MG

A075153 001 Feb 26, 1999

SUN PHARM INDS LTD 120MG

A077442 001 Sep 28, 2005

SUDAFED 24 HOUR

+! KENVUE BRANDS 240MG

N020021 002 Dec 15, 1992

PURIFIED WATER

SOLUTION;OPHTHALMIC

PUR-WASH

+! NIAGARA PHARMS 98.3%

N022305 001 Sep 01, 2011

RANITIDINE HYDROCHLORIDE

TABLET;ORAL

RANITIDINE HYDROCHLORIDE

DR REDDYS LABS LTD EQ 75MG BASE

A075294 001 Mar 28, 2000

EQ 150MG BASE

A078192 001 Aug 31, 2007

TERBINAFINE

GEL;TOPICAL

LAMISIL AT

+! KARO HLTHCARE 1%

N021958 001 Jul 24, 2006

TERBINAFINE HYDROCHLORIDE

CREAM;TOPICAL

LAMISIL

+! KARO HLTHCARE 1%

N020980 001 Mar 09, 1999

TERBINAFINE HYDROCHLORIDE

TARO 1%

A077511 001 Jul 02, 2007

SOLUTION;TOPICAL

LAMISIL AT

+! KARO HLTHCARE 1%

N021124 001 Mar 17, 2000

SPRAY;TOPICAL

LAMISIL AT

+! KARO HLTHCARE 1%

N021124 002 Mar 17, 2000

TIOCONAZOLE

OINTMENT;VAGINAL

TIOCONAZOLE

PERRIGO 6.5%

A075915 001 Nov 21, 2001

VAGISTAT-1

+! COMBE 6.5%

N020676 001 Feb 11, 1997

TRIAMCINOLONE ACETONIDE

SPRAY, METERED;NASAL

NASACORT ALLERGY 24 HOUR

+! CHATTEM SANOFI 0.055MG/SPRAY

N020468 002 Oct 11, 2013

TRIAMCINOLONE ACETONIDE

APOTEX 0.055MG/SPRAY

A214615 001 Jan 19, 2023

PERRIGO PHARMA INTL 0.055MG/SPRAY

A078104 002 Nov 14, 2014

**DRUG PRODUCTS WITH APPROVAL UNDER SECTION 505 OF THE ACT
ADMINISTERED BY THE CENTER FOR BIOLOGICS EVALUATION AND RESEARCH LIST**

0.9% SODIUM CHLORIDE INJECTION USP SOLUTION

SOLUTION; STORAGE/PROCESSING ONLY

NONE

FRESENIUS KABI AG

N125695

Sep 05, 2019

ANTICOAGULANT 4% SODIUM CITRATE SOLUTION USP

SOLUTION; STORAGE/PROCESSING ONLY

NONE

HAEMONETICS MANUFACTURING INC

N760305

Jun 30, 1978

ANTICOAGULANT CITRATE DEXTROSE SOLUTION (ACD)

SOLUTION; STORAGE/PROCESSING ONLY

NONE

CITRA LABS LLC

N020037

Aug 26, 2003

ACD-A SOLUTION

TERUMO BCT INC

A010228

Feb 25, 2002

ADSOL WITH ACD-A

FENWAL INC

N000922

Aug 29, 2002

ANTICOAGULANT CITRATE DEXTROSE SOLUTION FORMULA A

HAEMONETICS CORP

A980728

Feb 06, 2002

ADDITIVE SOLUTION 3; AS-3

TERUMO BCT INC

N001214

May 29, 2002

ANTICOAGULANT CITRATE DEXTROSE SOLUTION A (ACD-A)

TERUMO BCT INC

A110057

May 11, 2012

ANTICOAGULANT CITRATE DEXTROSE SOLUTION USP

SOLUTION; STORAGE/PROCESSING ONLY

NONE

FENWAL INC

N160918

Mar 17, 1978

ANTICOAGULANT CITRATE PHOSPHATE 2X DEXTROSE SOLUTION (CP2D)

SOLUTION; STORAGE/PROCESSING ONLY

CITRATE PHOSPHATE DOUBLE DEXTROSE/ADDITIVE SOLUTION 3
HAEMONETICS CORP

N000127

Mar 17, 1978

ANTICOAGULANT CITRATE PHOSPHATE DEXTROSE ADENINE SOLUTION

SOLUTION; STORAGE/PROCESSING ONLY

NONE

TERUMO MEDICAL CORP

N820528

Nov 03, 1982

ANTICOAGULANT CITRATE PHOSPHATE DEXTROSE ADENINE SOLUTION (CPDA)

SOLUTION; STORAGE/PROCESSING ONLY

CPDA-1 BLOOD-PACK UNIT (PL 146 PLASTIC)

N770420

May 12, 1978

FENWAL INC

**DRUG PRODUCTS WITH APPROVAL UNDER SECTION 505 OF THE ACT
ADMINISTERED BY THE CENTER FOR BIOLOGICS EVALUATION AND RESEARCH LIST**

ANTICOAGULANT CITRATE PHOSPHATE DEXTROSE ADENINE SOLUTION USP

SOLUTION; STORAGE/PROCESSING ONLY

BLOOD PACK UNIT CPDA-1 IN PLASTIC CONTAINER

FENWAL INC

N940404 Jul 28, 1994

ANTICOAGULANT CITRATE PHOSPHATE DEXTROSE ADENINE-1 SOLUTION

SOLUTION; STORAGE/PROCESSING ONLY

NONE

HAEMONETICS MANUFACTURING INC

N800077 Nov 06, 1980

ANTICOAGULANT CITRATE PHOSPHATE DEXTROSE SOLUTION

SOLUTION; STORAGE/PROCESSING ONLY

ADSOL IN PLASTIC CONTAINER

FENWAL INC

N900223 Dec 27, 1991

ANTICOAGULANT CITRATE PHOSPHATE DEXTROSE SOLUTION (CPD)

SOLUTION; STORAGE/PROCESSING ONLY

CPD ANTICOAGULANT IN PL 2209 PLASTIC CONTAINER

FENWAL INC

N900224 Dec 27, 1991

MACOPRODUCTIONS SAS CPD/AS-1: MACOPHARMA LEUCOFLEX MTL1
LEUKOREDUCTION SYSTEM FOR BLOOD COMPONENTS KNOWN AS MTL1-WB

MACOPHARMA

N040083 Nov 21, 2005

NONE

TERUMO BCT INC

A070025 Jan 06, 2009

ANTICOAGULANT CITRATE PHOSPHATE DEXTROSE SOLUTION USP

SOLUTION; STORAGE/PROCESSING ONLY

NONE

FENWAL INC

N170401 Dec 06, 1977

N811012 Jun 28, 1983

GLOBAL LIFE SCIENCE SOLUTIONS USA, LLC

N800222 Aug 23, 1982

TERUMO MEDICAL CORP

N781211 JUN 10, 1981

ANTICOAGULANT CITRATE PHOSPHATE DEXTROSE SOLUTION USP WITH: AS-1:
DEXTROSE USP; SODIUM CHLORIDE USP; MANNITOL USP; ADENINE

SOLUTION; STORAGE/PROCESSING ONLY

ADSOL RED BLOOD CELL PRESERVATIVE SOLUTION

FENWAL INC

N811104 May 16, 1983

ANTICOAGULANT CITRATE PHOSPHATE DEXTROSE SOLUTION USP WITH: AS-5:
DEXTROSE USP; SODIUM CHLORIDE USP; MANNITOL USP; ADENINE

SOLUTION; STORAGE/PROCESSING ONLY

OPTISOL RED BLOOD CELL PRESERVATIVE SOLUTION

TERUMO MEDICAL CORP

N880217 Oct 07, 1988

**DRUG PRODUCTS WITH APPROVAL UNDER SECTION 505 OF THE ACT
ADMINISTERED BY THE CENTER FOR BIOLOGICS EVALUATION AND RESEARCH LIST**

ANTICOAGULANT CITRATE PHOSPHATE DOUBLE DEXTROSE SOLUTION WITH:
AS-3: CITRIC ACID USP; MONOBASIC SODIUM PHOSPHATE USP; SODIUM
CHLORIDE USP; ADENINE; DEXTROSE USP; SODIUM CITRATE USP

SOLUTION; STORAGE/PROCESSING ONLY

AS-3 NUTRICEL ADDITIVE SYSTEM

0.042GM/100ML;0.276GM/100ML;
0.410GM/100ML;0.30GM/100ML;
1.10GM/100ML;0.588GM/100ML

HAEMONETICS MANUFACTURING INC

N820915

Oct 19, 1984

ANTICOAGULANT SODIUM CITRATE 4% SOLUTION

SOLUTION; STORAGE/PROCESSING ONLY

NONE

BAXTER HEALTHCARE CORPORATION

N980123

Mar 03, 2000

LABORATORIES GRIFOIS, S.A.

A125697

Oct 25, 2019

TERUMO BCT

A125608

Jun 26, 2018

CSL PLASMA INC

A125750

Apr 25, 2022

ANTICOAGULANT SODIUM CITRATE SOLUTION

SOLUTION; STORAGE/PROCESSING ONLY

TRICITRASOL

CITRA LABS, LLC

N010409

Jul 10, 2003

ANTICOAGULANT SODIUM CITRATE SOLUTION USP

SOLUTION; STORAGE/PROCESSING ONLY

NONE

FENWAL INC

N770923

Jan 20, 1978

TERUMO MEDICAL CORP

N781214

Feb 08, 1980

CORD BLOOD STERILE COLLECTION BAG, ANTICOAGULANT CITRATE
PHOSPHATE DEXTROSE SOLUTION (CPD)

SOLUTION; STORAGE/PROCESSING ONLY

NONE

MACOPHARMA

N125552

Dec 21, 2016

DEXTRAN 1 IN SODIUM CHLORIDE 0.6%

INJECTABLE; INTRAVENOUS

PROMIT

MEDA AB

N830715

Oct 30, 1984

DEXTRAN 40, 10% IN DEXTROSE 5%

INJECTABLE; INTRAVENOUS

LMD IN GLASS BOTTLE

10GM/100ML

HOSPIRA INC

A720563

Oct 30, 1992

**DRUG PRODUCTS WITH APPROVAL UNDER SECTION 505 OF THE ACT
ADMINISTERED BY THE CENTER FOR BIOLOGICS EVALUATION AND RESEARCH LIST**

DEXTRAN 40, 10% IN SODIUM CHLORIDE 0.9%

INJECTABLE; INTRAVENOUS

LMD IN PLASTIC CONTAINER 10GM/100ML;0.9GM/100ML

HOSPIRA INC

A720562 Oct 30, 1992

HETASTARCH 6% IN LACTATED ELECTROLYTE INJECTION

INJECTABLE; INTRAVENOUS

HEXTEND 6GM/100ML

BIOTIME INC

N200952 Mar 31, 1999

HETASTARCH 6% IN SODIUM CHLORIDE 0.9%

INJECTABLE; INTRAVENOUS

6% HETASTARCH IN SODIUM CHLORIDE

0.9% IN PLASTIC CONTAINER 6GM/100ML

HOSPIRA INC

A740193 Jan 30, 1995

HESPAN IN PLASTIC CONTAINER 6GM/100ML

B BRAUN MEDICAL INC

N890105 Apr 04, 1991

NONE 6GM/100ML

TEVA PHARMACEUTICALS USA INC

A740592 Nov 12, 1998

ISOPLATE SOLUTION IN THE 500 ML EXCEL CONTAINER

SOLUTION; STORAGE/PROCESSING ONLY

ISOPLATE SOLUTION

TERUMO BCT

N090067 Mar 05, 2013

LEUKOCYTE REDUCTION FILTRATION SYSTEM FOR WHOLE BLOOD WITH CPD
ANTICOAGULANT AND SOLX ADDITIVE

SOLUTION; STORAGE/PROCESSING ONLY

LEUKOSEP HWB-600-XL

HEMERUS MEDICAL, LLC

N110059 Apr 25, 2013

RED BLOOD CELL PROCESSING SOLUTION

SOLUTION; STORAGE/PROCESSING ONLY

REJUVESOL

CITRA LABS LLC

N950522 Feb 26, 1997

SODIUM CHLORIDE; SODIUM ACETATE; SODIUM CITRATE DIHYDRATE;
SODIUM PHOSPHATE, DIABASIC ANHYDROUS; SODIUM PHOSPHATE
MONOBASIC, MONOHYDRATE

SOLUTION; STORAGE/PROCESSING ONLY

INTERSOL SOLUTION

2.26GM/500ML; 2.21GM/500ML;
1.59GM/500ML; 1.53GM/500ML;
0.465GM/500ML

FRESENIUS KABI USA, LLC

N080041 Dec 09, 2009

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ABACAIVIR SULFATE

TABLET;ORAL

ABACAIVIR SULFATE

APOTEX INC

EQ 300MG BASE

A201570 001 Dec 17, 2012

ZIAGEN

+ VIIIV HLTHCARE

EQ 300MG BASE **

N020977 001 Dec 17, 1998

ABACAIVIR SULFATE; LAMIVUDINE

TABLET;ORAL

ABACAIVIR SULFATE AND LAMIVUDINE

AUROBINDO PHARMA

EQ 600MG BASE;300MG

A206151 001 Mar 28, 2017

TEVA PHARMS USA

EQ 600MG BASE;300MG

A079246 001 Sep 29, 2016

ZYDUS PHARMS

EQ 600MG BASE;300MG

A208990 001 Nov 15, 2018

EPZICOM

+ VIIIV HLTHCARE

EQ 600MG BASE;300MG **

N021652 001 Aug 02, 2004

TABLET, FOR SUSPENSION;ORAL

ABACAIVIR SULFATE AND LAMIVUDINE

+ NORVIUM BIOSCIENCE

EQ 60MG BASE;30MG

N204311 001 Dec 22, 2023

ABACAIVIR SULFATE; LAMIVUDINE; ZIDOVUDINE

TABLET;ORAL

TRIZIVIR

+ VIIIV HLTHCARE

EQ 300MG BASE;150MG;300MG **

N021205 001 Nov 14, 2000

ABAMETAPIR

LOTION;TOPICAL

XEGLYZE

+ HATCHTECH

0.74%

N206966 001 Jul 24, 2020

ABARELIX

INJECTABLE;INTRAMUSCULAR

PLENAXIS

SPECIALITY EUROPEAN

100MG/VIAL

N021320 001 Nov 25, 2003

ABIRATERONE ACETATE

TABLET;ORAL

ABIRATERONE ACETATE

TEVA PHARMS USA

125MG

A212206 001 Jun 24, 2022

ACAMPROSATE CALCIUM

TABLET, DELAYED RELEASE;ORAL

ACAMPROSATE CALCIUM

BARR LABS DIV TEVA

333MG

A200143 001 Nov 18, 2013

GLENMARK PHARMS LTD

333MG

A202229 001 Jul 16, 2013

CAMPRAL

+ FOREST LABS

333MG **

N021431 001 Jul 29, 2004

ACARBOSE

TABLET;ORAL

ACARBOSE

DASH PHARMS

25MG

A091053 001 Jan 06, 2011

50MG

A091053 002 Jan 06, 2011

100MG

A091053 003 Jan 06, 2011

HANGZHOU ZHONGMEI

25MG

A213821 001 Aug 18, 2020

50MG

A213821 002 Aug 18, 2020

100MG

A213821 003 Aug 18, 2020

PRECOSE

+ BAYER HLTHCARE

25MG **

N020482 004 May 29, 1997

+

50MG **

N020482 001 Sep 06, 1995

+

100MG **

N020482 002 Sep 06, 1995

ACEBUTOLOL HYDROCHLORIDE

CAPSULE;ORAL

ACEBUTOLOL HYDROCHLORIDE

NORVIUM BIOSCIENCE

EQ 200MG BASE

A074288 001 Apr 24, 1995

EQ 400MG BASE

A074288 002 Apr 24, 1995

SECTRAL

+ PROMIUS PHARMA

EQ 200MG BASE **

N018917 001 Dec 28, 1984

+

EQ 400MG BASE **

N018917 003 Dec 28, 1984

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ACETAMINOPHEN

INJECTABLE; INJECTION

INJECTAPAP

ORTHO MCNEIL PHARM 100MG/ML

N017785 001 Mar 07, 1986

POWDER; INTRAVENOUS

ACETAMINOPHEN

+ RISING 1GM/VIAL

N206610 001 Jan 15, 2021

SOLUTION; INTRAVENOUS

ACETAMINOPHEN

ZYDUS PHARMS 1GM/100ML (10MG/ML)

A216467 001 Oct 25, 2022

OFIRMEV

+ MALLINCKRODT HOSP 1GM/100ML (10MG/ML) **

N022450 001 Nov 02, 2010

SUPPOSITORY; RECTAL

ACEPHEN

COSETTE 120MG

A072218 001 Mar 27, 1992

120MG

N018060 001

325MG

A072344 001 Mar 27, 1992

325MG

N018060 003 Dec 18, 1986

650MG

A072237 001 Mar 27, 1992

650MG

N018060 002

ACETAMINOPHEN

ABLE 120MG

A073106 001 Feb 27, 1995

325MG

A073107 001 Feb 27, 1995

650MG

A073108 001 Feb 27, 1995

ACINO PRODS 120MG

A071010 001 May 12, 1987

650MG

A071011 001 May 12, 1987

+ TARO 650MG

N018337 001

NEOPAP

POLYMEDICA 120MG

N016401 001

TYLENOL

J AND J CONSUMER INC 120MG

N017756 002

650MG

N017756 001

TABLET, EXTENDED RELEASE; ORAL

ACETAMINOPHEN

SUN PHARM INDS LTD 650MG

A078569 001 Dec 14, 2011

650MG

A090205 001 Nov 18, 2009

ACETAMINOPHEN; ASPIRIN; CODEINE PHOSPHATE

CAPSULE; ORAL

ACETAMINOPHEN, ASPIRIN, AND CODEINE PHOSPHATE

MIKART 150MG; 180MG; 15MG

A081095 001 Oct 26, 1990

150MG; 180MG; 30MG

A081096 001 Oct 26, 1990

150MG; 180MG; 60MG

A081097 001 Oct 26, 1990

CODEINE, ASPIRIN, APAP FORMULA NO. 2

+ SCHERER LABS 150MG; 180MG; 15MG

A085640 001

CODEINE, ASPIRIN, APAP FORMULA NO. 3

+ SCHERER LABS 150MG; 180MG; 30MG

A085639 001

CODEINE, ASPIRIN, APAP FORMULA NO. 4

SCHERER LABS 150MG; 180MG; 60MG

A085638 001

ACETAMINOPHEN; BENZHYDROCODONE HYDROCHLORIDE

TABLET; ORAL

APADAZ

+ ZEVRA THERAP 325MG; EQ 4.08MG BASE

N208653 002 Jan 04, 2019

+ 325MG; EQ 6.12MG BASE

N208653 001 Feb 23, 2018

+ 325MG; EQ 8.16MG BASE

N208653 003 Jan 04, 2019

ACETAMINOPHEN; BUTALBITAL

CAPSULE; ORAL

BANCAP

FOREST PHARMS 325MG; 50MG

A088889 001 Jan 16, 1986

BUCET

MALLINCKRODT 650MG; 50MG

A088991 001 Jun 28, 1985

PHRENILIN FORTE

VALEANT 650MG; 50MG

A088831 001 Jun 19, 1985

TENCON

MALLINCKRODT 650MG; 50MG

A089405 001 May 15, 1990

TRIAPRIN

DUNHALL 325MG; 50MG

A089268 001 Jul 02, 1987

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ACETAMINOPHEN; BUTALBITAL

TABLET; ORAL

BUTALBITAL AND ACETAMINOPHEN

ALVOGEN	300MG;50MG	A207635 001	Jun 05, 2017
	325MG;50MG	A205120 001	Oct 30, 2015
HALSEY	325MG;50MG	A089568 001	Oct 05, 1988
+ WATSON LABS	325MG;50MG	A087550 001	Oct 19, 1984
BUTAPAP			
MIKART	650MG;50MG	A089988 001	Oct 26, 1992
PHRENILIN			
+ VALEANT	325MG;50MG **	A087811 001	Jun 19, 1985
SEDAPAP			
MAYRAND	650MG;50MG	A088944 001	Oct 17, 1985

ACETAMINOPHEN; BUTALBITAL; CAFFEINE

CAPSULE; ORAL

ANOQUAN

SHIRE	325MG;50MG;40MG	A087628 001	Oct 01, 1986
BUTALBITAL, ACETAMINOPHEN AND CAFFEINE			
+ GILBERT LABS	325MG;50MG;40MG **	A088825 001	Dec 05, 1984
GRAHAM DM	325MG;50MG;40MG	A088743 001	Jul 18, 1985
	325MG;50MG;40MG	A088765 001	Mar 27, 1985
	325MG;50MG;40MG	A089067 001	Apr 19, 1985
HIKMA	300MG;50MG;40MG	A215135 001	Mar 25, 2022
	325MG;50MG;40MG	A215135 002	Mar 25, 2022
	500MG;50MG;40MG	A040261 001	Oct 28, 1998
KEY THERAP	300MG;50MG;40MG	A206615 001	Aug 04, 2017
MALLINCKRODT	325MG;50MG;40MG	A088758 001	Mar 27, 1985
ESGIC-PLUS			
MIKART	500MG;50MG;40MG	A040085 001	Mar 28, 1996
FEMCET			
MALLINCKRODT	325MG;50MG;40MG	A089102 001	Jun 19, 1985
MEDIGESIC PLUS			
US CHEM	325MG;50MG;40MG	A089115 001	Jan 14, 1986
TRIAD			
MALLINCKRODT	325MG;50MG;40MG	A089023 001	Jun 19, 1985

TABLET; ORAL

BUTALBITAL, ACETAMINOPHEN AND CAFFEINE

ABLE	325MG;50MG;40MG	A040390 001	Jul 23, 2001
	500MG;50MG;40MG	A040394 001	Jul 23, 2001
ALVOGEN	325MG;50MG;40MG	A204984 001	Jan 10, 2017
GILBERT LABS	325MG;50MG;40MG	A087629 001	Nov 13, 1984
HIKMA	500MG;50MG;40MG	A040336 001	Aug 18, 1999
HIKMA PHARMS	325MG;50MG;40MG	A089718 001	Jun 12, 1995
MIKART	750MG;50MG;40MG	A040496 001	Dec 23, 2003
MIRROR PHARMS LLC	500MG;50MG;40MG	A040883 001	Dec 23, 2008
NESHER PHARMS	325MG;50MG;40MG	A211543 001	Jul 17, 2020
SPECGX LLC	325MG;50MG;40MG	A087804 001	Jan 24, 1985
SUN PHARM INDUSTRIES	325MG;50MG;40MG	A040601 001	Jul 29, 2005
VINTAGE PHARMS	500MG;50MG;40MG	A040513 001	Aug 25, 2003
WATSON LABS	325MG;50MG;40MG	A089536 001	Feb 16, 1988
	500MG;50MG;40MG	A040267 001	Jul 30, 1998
ESGIC			
FOREST PHARMS	325MG;50MG;40MG	A089660 001	Dec 23, 1988
ESGIC-PLUS			
MIKART	500MG;50MG;40MG	A089451 001	May 23, 1988

ACETAMINOPHEN; BUTALBITAL; CAFFEINE; CODEINE PHOSPHATE

CAPSULE; ORAL

BUTALBITAL, ACETAMINOPHEN, CAFFEINE AND CODEINE PHOSPHATE

ABLE	325MG;50MG;40MG;30MG	A076528 001	Aug 21, 2003
HIKMA INTL PHARMS	325MG;50MG;40MG;30MG	A075618 001	Mar 23, 2001
NOSTRUM LABS INC	325MG;50MG;40MG;30MG	A075929 001	Apr 22, 2002
PHRENILIN WITH CAFFEINE AND CODEINE			
VALEANT	325MG;50MG;40MG;30MG	A074911 001	Aug 22, 2001

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ACETAMINOPHEN; CAFFEINE; DIHYDROCODEINE BITARTRATE

CAPSULE;ORAL

ACETAMINOPHEN, CAFFEINE, AND DIHYDROCODEINE BITARTRATE

MIKART	356.4MG;30MG;16MG	A040109	001	Aug 26, 1997
WRASER PHARMS LLC	356.4MG;30MG;16MG	A040688	001	Apr 03, 2007
DHC PLUS				
PHARM RES ASSOC	356.4MG;30MG;16MG	A088584	001	Mar 04, 1986
SYNALGOS-DC-A				
LEITNER PHARMS	356.4MG;30MG;16MG	A089166	001	May 14, 1986
TREZIX				
KEY THERAP	320.5MG;30MG;16MG	A204785	001	Nov 26, 2014

TABLET;ORAL

ACETAMINOPHEN, CAFFEINE AND DIHYDROCODEINE BITARTRATE

LARKEN LABS INC	325MG;30MG;16MG	A204209	001	Sep 30, 2016
ACETAMINOPHEN, CAFFEINE, AND DIHYDROCODEINE BITARTRATE				
BOCA PHARMA LLC	712.8MG;60MG;32MG	A040701	001	Apr 03, 2007
MIKART	712.8MG;60MG;32MG	A040316	001	Apr 28, 1999
WEST-WARD PHARM CORP	712.8MG;60MG;32MG	A040637	001	Sep 22, 2006

ACETAMINOPHEN; CLEMASTINE FUMARATE; PSEUDOEPHEDRINE HYDROCHLORIDE

TABLET;ORAL

TAVIST ALLERGY/SINUS/HEADACHE

NOVARTIS	500MG;EQ 0.25MG BASE;30MG	N021082	001	Mar 01, 2001
----------	---------------------------	---------	-----	--------------

ACETAMINOPHEN; CODEINE PHOSPHATE

CAPSULE;ORAL

ACETAMINOPHEN AND CODEINE PHOSPHATE

TEVA	300MG;15MG	A088537	001	Jun 04, 1984
	300MG;30MG	A088324	001	Dec 29, 1983
	300MG;60MG	A088599	001	Jun 01, 1984
PHENAPHEN W/ CODEINE NO. 2				
ROBINS AH	325MG;15MG	A084444	001	
PHENAPHEN W/ CODEINE NO. 3				
+ ROBINS AH	325MG;30MG	A084445	001	
PHENAPHEN W/ CODEINE NO. 4				
+ ROBINS AH	325MG;60MG	A084446	001	
PROVAL #3				
SOLVAY	325MG;30MG	A085685	001	
TYLENOL W/ CODEINE NO. 3				
ORTHO MCNEIL PHARM	300MG;30MG	A087422	001	
TYLENOL W/ CODEINE NO. 4				
ORTHO MCNEIL PHARM	300MG;60MG	A087421	001	

SOLUTION;ORAL

ACETAMINOPHEN AND CODEINE PHOSPHATE

+ ACTAVIS MID ATLANTIC	120MG/5ML;12MG/5ML	A085861	001	
CHARTWELL MOLECULAR	120MG/5ML;12MG/5ML	A091238	001	Nov 10, 2011
DAVA PHARMS INC	120MG/5ML;12MG/5ML	A040098	001	Sep 20, 1996
PAI HOLDINGS	120MG/5ML;12MG/5ML	A040119	001	Apr 26, 1996
STRIDES PHARMA	120MG/5ML;12MG/5ML	A086366	001	
WOCKHARDT BIO AG	120MG/5ML;12MG/5ML	A087006	001	
TYLENOL W/ CODEINE				
+ ORTHO MCNEIL PHARM	120MG/5ML;12MG/5ML	A085057	001	

SUSPENSION;ORAL

CAPITAL AND CODEINE

ACTAVIS MID ATLANTIC	120MG/5ML;12MG/5ML	A085883	001	
VALEANT PHARMS LLC	120MG/5ML;12MG/5ML	A086024	001	

TABLET;ORAL

ACETAMINOPHEN AND CODEINE PHOSPHATE

ABLE	300MG;30MG	A040452	001	Aug 01, 2002
	300MG;60MG	A040459	001	Aug 01, 2002
AM THERAP				
	300MG;15MG	A089478	001	Mar 03, 1987
	300MG;15MG	A089481	001	Mar 03, 1987
	300MG;30MG	A089479	001	Mar 03, 1987
	300MG;30MG	A089482	001	Mar 03, 1987
	300MG;60MG	A089480	001	Mar 03, 1987
	300MG;60MG	A089483	001	Mar 03, 1987
CHARTWELL				
	300MG;15MG	A089997	001	Dec 28, 1994
	300MG;30MG	A089998	001	Dec 28, 1994
	300MG;60MG	A089999	001	Dec 28, 1994
DURAMED PHARMS BARR	300MG;15MG	A040223	001	Nov 18, 1997

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ACETAMINOPHEN; CODEINE PHOSPHATE

TABLET;ORAL

ACETAMINOPHEN AND CODEINE PHOSPHATE

	300MG;15MG	A088353	001	Feb 06, 1984
	300MG;30MG	A040223	002	Nov 18, 1997
	300MG;30MG	A088354	001	Feb 06, 1984
	300MG;60MG	A040223	003	Nov 18, 1997
	300MG;60MG	A088355	001	Feb 06, 1984
EVERYLIFE	325MG;30MG	A085217	001	
FOSUN PHARMA	300MG;30MG	A081250	001	Jul 16, 1992
	300MG;60MG	A081249	001	Jul 16, 1992
HALSEY	300MG;15MG	A083871	001	
	300MG;30MG	A083872	001	
	300MG;60MG	A086549	001	
KV PHARM	300MG;30MG	A085288	001	
	300MG;60MG	A085365	001	
	325MG;15MG	A085364	001	
	325MG;45MG **	A085363	001	
LEDERLE	300MG;30MG	A087141	001	
MIKART	300MG;30MG	A089238	001	Feb 25, 1986
	300MG;60MG	A089244	001	Feb 25, 1986
	650MG;30MG	A089231	001	Mar 03, 1986
	650MG;60MG	A089363	001	Sep 09, 1991
MUTUAL PHARM	300MG;15MG	A085795	001	
	300MG;30MG	A085794	001	
	300MG;60MG	A087653	001	Apr 13, 1982
NOSTRUM LABS INC	300MG;15MG	A088629	002	Mar 06, 1985
	300MG;30MG	A088629	003	Mar 06, 1985
+	300MG;60MG	A088629	001	Mar 06, 1985
PURACAP PHARM	300MG;30MG	A087762	001	Dec 10, 1982
+	300MG;30MG	A086681	001	
	300MG;30MG	A089080	001	Jul 17, 1986
	300MG;60MG	A086683	001	
RHODES PHARMS	300MG;15MG	A089673	002	Feb 10, 1988
	300MG;30MG	A089673	003	Feb 10, 1988
	300MG;60MG	A089673	001	Feb 10, 1988
ROXANE	300MG;15MG	A084659	001	
	300MG;30MG	A084656	001	
	300MG;60MG	A084667	001	
	500MG;15MG	A089511	001	Apr 25, 1989
	500MG;30MG	A089512	001	Apr 25, 1989
	500MG;60MG	A089513	001	Apr 25, 1989
+	SANDOZ	300MG;15MG	A087433	001
	300MG;30MG	A085291	002	
+	300MG;30MG	A085917	001	
	300MG;60MG	A085964	001	
	300MG;60MG	A087423	001	
SUN PHARM INDS LTD	300MG;30MG	A085868	001	
SUPERPHARM	300MG;15MG	A089183	001	Oct 18, 1985
	300MG;30MG	A089184	001	Oct 18, 1985
	300MG;30MG	A089253	001	May 19, 1986
	300MG;60MG	A089185	001	Oct 18, 1985
	300MG;60MG	A089254	001	May 19, 1986
USL PHARMA	300MG;30MG	A087919	001	Jun 22, 1982
	300MG;60MG	A087920	001	Jun 22, 1982
VALEANT PHARM INTL	300MG;30MG	A085896	001	
VITARINE	300MG;30MG	A085676	001	
WARNER CHILCOTT	300MG;15MG	A085992	001	
	300MG;30MG	A085218	002	
	300MG;60MG	A087306	001	
WATSON LABS	300MG;15MG	A087277	001	May 26, 1982
	300MG;30MG	A087276	001	May 26, 1982
	300MG;60MG	A087275	001	May 26, 1982
WATSON LABS FLORIDA	300MG;15MG	A040443	001	Jan 22, 2003
	300MG;30MG	A040443	002	Jan 22, 2003
	300MG;60MG	A040443	003	Jan 22, 2003
WHITEWORTH TOWN PLSN	300MG;30MG	A084360	001	
	300MG;60MG	A085607	001	

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ACETAMINOPHEN; CODEINE PHOSPHATE

TABLET; ORAL

CAPITAL AND CODEINE

CARNRICK 325MG; 30MG A083643 001

CODRIX

WATSON LABS FLORIDA 500MG; 15MG A040447 001 Feb 26, 2003

500MG; 30MG A040441 001 Mar 27, 2003

500MG; 60MG A040488 001 Mar 28, 2003

EMPRACET W/ CODEINE PHOSPHATE #3

GLAXOSMITHKLINE 300MG; 30MG A083951 001

EMPRACET W/ CODEINE PHOSPHATE #4

GLAXOSMITHKLINE 300MG; 60MG A083951 002

PAPA-DEINE #3

VANGARD 300MG; 30MG A088037 001 Mar 20, 1984

PAPA-DEINE #4

VANGARD 300MG; 60MG A088715 001 Mar 20, 1984

PHENAPHEN-650 W/ CODEINE

ROBINS AH 650MG; 30MG A085856 001

TYLENOL W/ CODEINE

ORTHO MCNEIL PHARM 325MG; 7.5MG ** A085056 001

325MG; 15MG ** A085056 002

325MG; 30MG ** A085056 003

325MG; 60MG ** A085056 004

TYLENOL W/ CODEINE NO. 1

JANSSEN PHARMS 300MG; 7.5MG A085055 001

TYLENOL W/ CODEINE NO. 2

+ JANSSEN PHARMS 300MG; 15MG A085055 002

TYLENOL W/ CODEINE NO. 3

+ JANSSEN PHARMS 300MG; 30MG A085055 003

TYLENOL W/ CODEINE NO. 4

+ JANSSEN PHARMS 300MG; 60MG A085055 004

ACETAMINOPHEN; DEXBROMPHENIRAMINE MALEATE; PSEUDOEPHEDRINE SULFATE

TABLET, EXTENDED RELEASE; ORAL

DRIXORAL PLUS

SCHERING PLOUGH 500MG; 3MG; 60MG N019453 001 May 22, 1987

ACETAMINOPHEN; HYDROCODONE BITARTRATE

CAPSULE; ORAL

ACETAMINOPHEN AND HYDROCODONE BITARTRATE

CENT PHARMS 500MG; 5MG A088898 001 Mar 27, 1985

ALLAY

IVAX PHARMS 500MG; 5MG A089907 001 Jan 13, 1989

BANCAP HC

FOREST PHARMS 500MG; 5MG A087961 001 Mar 17, 1983

CO-GESIC

CENT PHARMS 500MG; 5MG A089360 001 Mar 02, 1988

HYDROCODONE BITARTRATE AND ACETAMINOPHEN

MALLINCKRODT 500MG; 5MG A088956 001 Jul 19, 1985

500MG; 5MG A089006 001 Aug 09, 1985

MIKART 500MG; 5MG A081067 001 Nov 30, 1989

500MG; 5MG A081068 001 Nov 30, 1989

500MG; 5MG A081069 001 Nov 30, 1989

500MG; 5MG A081070 001 Nov 30, 1989

500MG; 5MG A089008 001 Feb 21, 1986

LORCET-HD

MALLINCKRODT 500MG; 5MG A087336 001 Jul 08, 1982

SOLUTION; ORAL

HYDROCODONE BITARTRATE AND ACETAMINOPHEN

GENUS 325MG/15ML; 7.5MG/15ML A200343 001 Jan 25, 2012

MALLINCKRODT 500MG/15ML; 7.5MG/15ML A040418 001 Jun 27, 2001

MALLINCKRODT INC 500MG/15ML; 10MG/15ML A040508 001 Aug 29, 2003

MIKART 500MG/15ML; 5MG/15ML A081226 001 Oct 27, 1992

500MG/15ML; 5MG/15ML A089557 001 Apr 29, 1992

500MG/15ML; 7.5MG/15ML A081051 001 Aug 28, 1992

NESHER PHARMS 500MG/15ML; 7.5MG/15ML A040366 001 Jan 23, 2002

PHARM ASSOC 500MG/15ML; 7.5MG/15ML A040182 001 Mar 13, 1998

TRIS PHARMA INC 300MG/15ML; 10MG/15ML A201295 002 Dec 30, 2021

325MG/15ML; 7.5MG/15ML A201295 001 Dec 30, 2021

VINTAGE PHARMS 500MG/15ML; 7.5MG/15ML A040520 001 Oct 30, 2003

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ACETAMINOPHEN; HYDROCODONE BITARTRATE

SOLUTION; ORAL

ZYFREL

CYPRESS PHARM INC 325MG/15ML; 7.5MG/15ML A090468 001 Apr 14, 2016

TABLET; ORAL

ANEXSIA

MALLINCKRODT 500MG; 5MG A089160 001 Apr 23, 1987

750MG; 10MG A040468 001 Oct 31, 2002

ANEXSIA 7.5/650

MALLINCKRODT 650MG; 7.5MG A089725 001 Sep 30, 1987

CO-GESIC

UCB INC 500MG; 5MG A087757 001 May 03, 1982

DURADYNE DHC

FOREST PHARMS 500MG; 5MG A087809 001 Mar 17, 1983

HY-PHEN

ASCHEP 500MG; 5MG A087677 001 May 03, 1982

HYDROCODONE BITARTRATE AND ACETAMINOPHEN

ABLE 325MG; 5MG A040478 001 Nov 08, 2002

325MG; 7.5MG A040464 001 Oct 23, 2002

325MG; 10MG A040464 002 Oct 23, 2002

500MG; 5MG A040477 001 Nov 06, 2002

500MG; 7.5MG A040490 001 May 21, 2003

500MG; 10MG A040473 001 Nov 06, 2002

650MG; 7.5MG A040474 001 Jan 02, 2003

650MG; 10MG A040476 001 Oct 23, 2002

750MG; 7.5MG A040469 001 Oct 25, 2002

ACTAVIS LABS FL INC 300MG; 5MG A206470 001 Jun 02, 2016

300MG; 7.5MG A206470 002 Jun 02, 2016

300MG; 10MG A206470 003 Jun 02, 2016

ALVOGEN 300MG; 5MG A208540 001 Nov 08, 2018

300MG; 7.5MG A208540 002 Nov 08, 2018

300MG; 10MG A208540 003 Nov 08, 2018

325MG; 2.5MG A209958 001 Oct 24, 2018

325MG; 5MG A209958 002 Oct 24, 2018

325MG; 7.5MG A209958 003 Oct 24, 2018

325MG; 10MG A209958 004 Oct 24, 2018

AMNEAL PHARMS 300MG; 10MG A207137 001 Nov 29, 2016

AMNEAL PHARMS NY 300MG; 5MG A206869 001 Jun 23, 2017

500MG; 5MG A040729 001 Aug 25, 2006

500MG; 7.5MG A040748 001 Aug 25, 2006

500MG; 10MG A040813 001 Feb 23, 2007

650MG; 7.5MG A040754 001 Aug 25, 2006

650MG; 10MG A040757 001 Aug 25, 2006

750MG; 7.5MG A040769 001 Aug 28, 2006

APIL 500MG; 10MG A040148 002 Feb 14, 1997

BARR 500MG; 2.5MG A040307 001 Jul 26, 2000

500MG; 5MG A040308 001 Jul 26, 2000

500MG; 5MG A088577 001 Dec 21, 1984

500MG; 7.5MG A040307 002 Jul 26, 2000

500MG; 10MG A040309 001 Jul 26, 2000

650MG; 7.5MG A040307 003 Jul 26, 2000

650MG; 10MG A040307 004 Jul 26, 2000

750MG; 7.5MG A040308 002 Jul 26, 2000

CARACO 500MG; 5MG A090265 001 Dec 23, 2008

500MG; 7.5MG A090265 002 Dec 23, 2008

500MG; 10MG A090265 003 Dec 23, 2008

650MG; 7.5MG A090380 001 Dec 23, 2008

650MG; 10MG A090380 002 Dec 23, 2008

660MG; 10MG A090380 003 Dec 23, 2008

750MG; 7.5MG A090380 004 Dec 23, 2008

CHARTWELL 325MG; 7.5MG A040248 001 Apr 28, 2000

325MG; 10MG A040248 002 Apr 28, 2000

ENDO OPERATIONS 300MG; 5MG A090415 001 Jan 24, 2011

300MG; 7.5MG A090415 002 Jan 24, 2011

300MG; 10MG A090415 003 Jan 24, 2011

HALSEY 500MG; 5MG A089554 001 Jun 12, 1987

IVAX PHARMS 500MG; 5MG A089696 001 Apr 21, 1988

LANNETT CO INC 300MG; 5MG A207171 001 Jun 20, 2017

300MG; 7.5MG A207171 002 Jun 20, 2017

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ACETAMINOPHEN; HYDROCODONE BITARTRATE

TABLET; ORAL

HYDROCODONE BITARTRATE AND ACETAMINOPHEN

	300MG;10MG	A207171	003	Jun 20, 2017
	325MG;5MG	A207172	001	Jun 22, 2017
	325MG;7.5MG	A207172	002	Jun 22, 2017
	325MG;10MG	A207172	003	Jun 22, 2017
MALLINCKRODT	500MG;5MG	A040084	002	Jun 01, 1995
	500MG;7.5MG	A040201	001	Feb 27, 1998
	500MG;10MG	A040201	002	Feb 27, 1998
	650MG;10MG	A040084	004	Oct 16, 1996
	660MG;10MG	A040084	003	Jul 29, 1996
	750MG;7.5MG	A040084	001	Jun 01, 1995
MIKART	325MG;2.5MG	A040846	001	Jun 09, 2010
	500MG;2.5MG	A089698	001	Aug 25, 1989
	500MG;5MG	A089271	001	Jul 16, 1986
	500MG;5MG	A089697	001	Jan 28, 1992
	500MG;7.5MG	A089699	001	Aug 25, 1989
	650MG;5MG	A040849	001	Jun 09, 2010
	650MG;7.5MG	A089689	001	Jun 29, 1988
	650MG;10MG	A081223	001	May 29, 1992
MUTUAL PHARM	500MG;5MG	A040236	001	Sep 25, 1997
	650MG;7.5MG	A040240	002	Nov 26, 1997
	650MG;10MG	A040240	001	Nov 26, 1997
	750MG;7.5MG	A040236	002	Sep 25, 1997
NOVEL LABS INC	300MG;5MG	A206142	001	Nov 14, 2016
	300MG;7.5MG	A206142	002	Nov 14, 2016
	300MG;10MG	A206142	003	Nov 14, 2016
	325MG;5MG	A206245	001	Dec 01, 2016
	325MG;7.5MG	A206245	002	Dec 01, 2016
	325MG;10MG	A206245	003	Dec 01, 2016
PRINSTON INC	325MG;5MG	A214928	001	Dec 30, 2021
	325MG;7.5MG	A214928	002	Dec 30, 2021
	325MG;10MG	A214928	003	Dec 30, 2021
RANBAXY	500MG;5MG	A040825	001	Aug 16, 2007
	500MG;10MG	A040824	001	Aug 16, 2007
RANBAXY LABS LTD	750MG;7.5MG	A040822	001	Aug 16, 2007
SANALUZ	325MG;5MG	A211690	001	Feb 07, 2020
	325MG;7.5MG	A211690	002	Feb 07, 2020
	325MG;10MG	A211690	003	Feb 07, 2020
SANDOZ	500MG;5MG	A040149	001	Jan 27, 1997
	750MG;7.5MG	A040149	002	Jan 27, 1997
STRIDES PHARMA	300MG;5MG	A205001	001	Jul 05, 2016
	300MG;7.5MG	A205001	002	Jul 05, 2016
	300MG;10MG	A205001	003	Jul 05, 2016
SUN PHARM INDS INC	325MG;5MG	A090118	001	Dec 23, 2008
	325MG;7.5MG	A090118	002	Dec 23, 2008
	325MG;10MG	A090118	003	Dec 23, 2008
SUN PHARM INDS LTD	325MG;10MG	A040826	001	Aug 16, 2007
UCB INC	500MG;10MG	A040210	001	Aug 13, 1997
	650MG;7.5MG	A040134	001	Nov 21, 1996
UPSHER SMITH LABS	325MG;5MG	A206484	001	Mar 24, 2017
	325MG;7.5MG	A206484	002	Mar 24, 2017
	325MG;10MG	A206484	003	Mar 24, 2017
USL PHARMA	500MG;5MG	A089290	001	May 29, 1987
	500MG;5MG	A089291	001	May 29, 1987
VINTAGE PHARMS	500MG;2.5MG	A040144	002	Apr 25, 1997
	500MG;5MG	A089831	001	Sep 07, 1988
	500MG;5MG	A089971	001	Dec 02, 1988
	500MG;7.5MG	A040144	001	Feb 22, 1996
	500MG;10MG	A040356	001	May 31, 2000
	650MG;7.5MG	A040155	001	Apr 14, 1997
	650MG;10MG	A040143	001	Feb 22, 1996
	660MG;10MG	A040358	001	May 31, 2000
	750MG;7.5MG	A040157	001	Apr 12, 1996
VINTAGE PHARMS LLC	500MG;5MG	A040281	001	Sep 30, 1998
	500MG;7.5MG	A040280	001	Sep 30, 1998
	650MG;7.5MG	A040280	002	Sep 30, 1998
	650MG;10MG	A040280	003	Sep 30, 1998

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ACETAMINOPHEN; HYDROCODONE BITARTRATE

TABLET; ORAL

HYDROCODONE BITARTRATE AND ACETAMINOPHEN

	750MG; 7.5MG	A040281	002	Sep 30, 1998
WATSON LABS	500MG; 2.5MG	A040123	003	Mar 04, 1996
	500MG; 2.5MG	A081079	001	Aug 30, 1991
	500MG; 5MG	A040122	001	Mar 04, 1996
	500MG; 5MG	A089883	001	Dec 01, 1988
	500MG; 7.5MG	A040123	004	Mar 04, 1996
	500MG; 7.5MG	A081080	001	Aug 30, 1991
	650MG; 7.5MG	A040094	001	Sep 29, 1995
	650MG; 7.5MG	A040123	001	Mar 04, 1996
	650MG; 10MG	A040094	002	Sep 29, 1995
	650MG; 10MG	A040123	002	Mar 04, 1996
	660MG; 10MG	A040094	003	Aug 08, 2000
	750MG; 7.5MG	A040122	002	Mar 04, 1996
	750MG; 7.5MG	A081083	001	Aug 30, 1991
	750MG; 10MG	A040094	004	Mar 22, 1999
WATSON LABS FLORIDA	500MG; 5MG	A040493	001	May 28, 2003
	660MG; 10MG	A040495	001	May 28, 2003
	750MG; 7.5MG	A040494	001	May 28, 2003
LORTAB				
UCB INC	500MG; 5MG	A087722	001	Jul 09, 1982
	500MG; 10MG	A040100	001	Jan 26, 1996
NORCET				
ABANA	500MG; 5MG	A088871	001	May 15, 1986
NORCO				
APIL	325MG; 2.5MG	A040148	004	Jul 07, 2014
	325MG; 5MG	A040099	001	Jun 25, 1997
	325MG; 5MG	A040148	005	Jul 07, 2014
	325MG; 7.5MG	A040148	003	Sep 12, 2000
	325MG; 10MG	A040148	001	Feb 14, 1997
TYCOLET				
ORTHO MCNEIL PHARM	500MG; 5MG	A089385	001	Aug 27, 1986
VICODIN				
ABBOTT	500MG; 5MG	A085667	001	
ABBVIE	500MG; 5MG	A088058	001	Jan 07, 1983
VICODIN ES				
ABBVIE	750MG; 7.5MG	A089736	001	Dec 09, 1988
VICODIN HP				
ABBVIE	660MG; 10MG	A040117	001	Sep 23, 1996
ZYDONE				
VINTAGE PHARMS LLC	400MG; 5MG	A040288	001	Nov 27, 1998
	400MG; 7.5MG	A040288	002	Nov 27, 1998
	400MG; 10MG	A040288	003	Nov 27, 1998

ACETAMINOPHEN; OXYCODONE HYDROCHLORIDE

CAPSULE; ORAL

OXYCODONE AND ACETAMINOPHEN

ACTAVIS ELIZABETH	500MG; 5MG	A040199	001	Dec 30, 1998
BARR	500MG; 5MG	A040304	001	Oct 02, 2000
DURAMED PHARMS BARR	500MG; 5MG	A040289	001	Mar 16, 1999
HALSEY	500MG; 5MG	A089994	001	May 04, 1989
MALLINCKRODT	500MG; 5MG	A040257	001	Aug 04, 1998
MUTUAL PHARM	500MG; 5MG	A040219	001	Jan 22, 1998
VINTAGE PHARMS	500MG; 5MG	A040106	001	Jul 30, 1996
VINTAGE PHARMS LLC	500MG; 5MG	A040303	001	Dec 30, 1999
WATSON LABS	500MG; 5MG	A040234	001	Oct 30, 1997
ROXILOX				
ROXANE	500MG; 5MG	A040061	001	Jul 03, 1995
TYLOX				
JANSSEN PHARMS	500MG; 5MG	A088790	001	Dec 12, 1984
TYLOX-325				
ORTHO MCNEIL PHARM	325MG; 5MG	A088246	001	Nov 08, 1984
SOLUTION; ORAL				
OXYCODONE AND ACETAMINOPHEN				
SPECGX LLC	325MG/5ML; 5MG/5ML	A040680	001	Sep 29, 2006
OXYCODONE HYDROCHLORIDE AND ACETAMINOPHEN				
ENDO OPERATIONS	325MG/5ML; 5MG/5ML	A203573	001	Dec 18, 2014

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ACETAMINOPHEN; OXYCODONE HYDROCHLORIDE

SOLUTION; ORAL

ROXICET

HIKMA 325MG/5ML; 5MG/5ML ** A089351 001 Dec 03, 1986

TABLET; ORAL

OXYCODONE 2.5/APAP 500

BRISTOL MYERS SQUIBB 500MG; 2.5MG A085910 001

OXYCODONE 5/APAP 500

BRISTOL MYERS SQUIBB 500MG; 5MG A085911 001

OXYCODONE AND ACETAMINOPHEN

ACTAVIS ELIZABETH 325MG; 5MG A040203 001 Mar 15, 1999

325MG; 7.5MG A040800 001 Apr 03, 2012

325MG; 10MG A040800 002 Apr 03, 2012

AMNEAL PHARMS NY 500MG; 7.5MG A040789 001 Nov 27, 2007

650MG; 10MG A040789 002 Nov 27, 2007

BARR 325MG; 5MG A087406 001

DR REDDYS LABS SA 325MG; 2.5MG A090177 001 Oct 20, 2008

325MG; 5MG A090177 002 Oct 20, 2008

325MG; 7.5MG A090177 003 Oct 20, 2008

325MG; 10MG A090177 004 Oct 20, 2008

500MG; 7.5MG A090177 005 Oct 20, 2008

650MG; 10MG A090177 006 Oct 20, 2008

DURAMED PHARMS BARR 325MG; 5MG A040272 001 Jun 30, 1998

ENDO OPERATIONS 325MG; 2.5MG A090733 001 Jul 11, 2013

325MG; 5MG A040105 001 Jul 30, 1996

325MG; 7.5MG A090734 001 Jul 11, 2013

325MG; 10MG A090734 002 Jul 11, 2013

LANNETT CO INC 325MG; 5MG A207333 001 Sep 25, 2017

325MG; 10MG A207333 002 Sep 25, 2017

MALLINCKRODT 500MG; 7.5MG A040550 001 Jun 30, 2004

650MG; 10MG A040550 002 Jun 30, 2004

MIKART 400MG; 2.5MG A040679 001 May 16, 2006

400MG; 5MG A040687 001 Apr 27, 2006

400MG; 7.5MG A040698 001 Apr 27, 2006

400MG; 10MG A040692 001 Apr 27, 2006

500MG; 10MG A040676 001 Apr 19, 2006

NESHER PHARMS 325MG; 2.5MG A210079 001 Dec 28, 2017

325MG; 5MG A210079 002 Dec 28, 2017

325MG; 7.5MG A210079 003 Dec 28, 2017

325MG; 10MG A210079 004 Dec 28, 2017

SANALUZ 325MG; 5MG A207574 001 Dec 13, 2016

SUN PHARM INDS INC 325MG; 2.5MG A090535 001 Dec 26, 2013

325MG; 5MG A090535 002 Dec 26, 2013

325MG; 7.5MG A090535 003 Dec 26, 2013

325MG; 10MG A090535 004 Dec 26, 2013

WATSON LABS 325MG; 5MG A040171 001 Oct 30, 1997

325MG; 7.5MG A040535 001 Sep 05, 2003

325MG; 10MG A040535 002 Sep 05, 2003

500MG; 7.5MG A040371 001 Dec 29, 2000

650MG; 10MG A040371 002 Dec 29, 2000

PERCOCET

+ ENDO OPERATIONS 325MG; 5MG ** A085106 002

VINTAGE PHARMS LLC 500MG; 7.5MG A040341 001 Jul 26, 1999

650MG; 10MG A040341 002 Jul 26, 1999

ROXICET

+ HIKMA 325MG; 5MG A087003 001

ROXICET 5/500

ROXANE 500MG; 5MG A089775 001 Jan 12, 1989

TABLET, EXTENDED RELEASE; ORAL

XARTEMIS XR

+ MALLINCKRODT INC 325MG; 7.5MG ** N204031 001 Mar 11, 2014

ACETAMINOPHEN; OXYCODONE HYDROCHLORIDE; OXYCODONE TEREPHTHALATE

CAPSULE; ORAL

TYLOX

ORTHO MCNEIL PHARM 500MG; 4.5MG; 0.38MG A085375 001

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ACETAMINOPHEN; PENTAZOCINE HYDROCHLORIDE

TABLET; ORAL

ACETAMINOPHEN AND PENTAZOCINE HYDROCHLORIDE

GAVIS PHARMS	650MG;EQ 25MG BASE	A076202	001	Aug 02, 2002
WATSON LABS	650MG;EQ 25MG BASE	A074699	001	Mar 24, 2000
TALACEN				
SANOFI AVENTIS US	650MG;EQ 25MG BASE	N018458	001	Sep 23, 1982

ACETAMINOPHEN; PROPOXYPHENE HYDROCHLORIDE

TABLET; ORAL

DARVOCET

AAIPHARMA LLC	325MG;32.5MG	N016844	001	
DOLENE AP-65				
LEDERLE	650MG;65MG	A085100	001	
PROPOXYPHENE HYDROCHLORIDE AND ACETAMINOPHEN				
MYLAN	325MG;32MG	A083689	001	
	650MG;65MG	A083978	001	
SANDOZ	650MG;65MG	A089959	001	Jul 18, 1989
VINTAGE PHARMS	650MG;65MG	A040507	001	Jul 30, 2003
WATSON LABS	650MG;65MG	A040139	001	Dec 16, 1996
WYGESIC				
CARACO	650MG;65MG	A084999	001	

ACETAMINOPHEN; PROPOXYPHENE NAPSYLATE

TABLET; ORAL

DARVOCET A500

XANODYNE PHARM	500MG;100MG	A076429	001	Sep 10, 2003
DARVOCET-N 100				
XANODYNE PHARM	650MG;100MG	N017122	002	
DARVOCET-N 50				
XANODYNE PHARM	325MG;50MG	N017122	001	
PROPACET 100				
TEVA	650MG;100MG	A070107	001	Jun 12, 1985
PROPOXYPHENE NAPSYLATE AND ACETAMINOPHEN				
ABLE	650MG;100MG	A075838	001	Jul 11, 2001
ACTAVIS ELIZABETH	650MG;100MG	A070910	001	Jan 02, 1987
CORNERSTONE	325MG;100MG	A076743	001	May 07, 2004
	500MG;100MG	A076750	001	Jun 28, 2004
HALSEY	325MG;50MG	A072105	001	May 13, 1988
	650MG;100MG	A072106	001	May 13, 1988
IVAX SUB TEVA PHARMS	650MG;100MG	A070146	001	Aug 02, 1985
MALLINCKRODT	650MG;100MG	A075738	001	Feb 02, 2001
MIRROR PHARMS	650MG;100MG	A077821	001	Feb 11, 2008
MUTUAL PHARM	325MG;50MG	A070115	001	Jun 12, 1985
	650MG;100MG	A070116	001	Jun 12, 1985
	650MG;100MG	A070615	001	Mar 21, 1986
	650MG;100MG	A070771	001	Mar 21, 1986
	650MG;100MG	A070775	001	Mar 21, 1986
MYLAN	650MG;100MG	A072195	001	Feb 16, 1988
MYLAN PHARMS INC	650MG;100MG	A070145	001	Jun 12, 1985
SANDOZ	650MG;100MG	A070443	001	Jan 23, 1986
SUPERPHARM	650MG;100MG	A071319	001	Jan 06, 1987
TEVA	650MG;100MG	A070732	001	Jan 03, 1986
	650MG;100MG	A074119	001	Dec 19, 1994
VINTAGE PHARMS	325MG;50MG	A074843	002	Feb 15, 2001
	650MG;100MG	A074843	001	Feb 12, 1997
WATSON LABS	325MG;50MG	A070398	001	Dec 18, 1986
	650MG;100MG	A070399	001	Dec 18, 1986
WATSON LABS FLORIDA	500MG;100MG	A077196	001	Jun 28, 2005
	650MG;100MG	A076609	001	Nov 16, 2004
WOCKHARDT LTD	325MG;50MG	A077677	001	Mar 16, 2007
	650MG;100MG	A077677	002	Mar 16, 2007

ACETAMINOPHEN; TRAMADOL HYDROCHLORIDE

TABLET; ORAL

TRAMADOL HYDROCHLORIDE AND ACETAMINOPHEN

CHARTWELL RX	325MG;37.5MG	A076475	001	Apr 21, 2005
GRAVITI PHARMS	325MG;37.5MG	A076914	001	Jul 26, 2006
MACLEODS PHARMS LTD	325MG;37.5MG	A206885	001	May 02, 2017
NOSTRUM LABS INC	325MG;37.5MG	A078778	001	Apr 07, 2014
SUN PHARM INDS INC	325MG;37.5MG	A077184	001	Dec 16, 2005

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ACETAMINOPHEN; TRAMADOL HYDROCHLORIDE

TABLET; ORAL

ULTRACET

+ JANSSEN PHARMS 325MG; 37.5MG N021123 001 Aug 15, 2001

ACETAZOLAMIDE

CAPSULE, EXTENDED RELEASE; ORAL

ACETAZOLAMIDE

RISING 500MG A203917 001 Jun 18, 2019

DIAMOX

+ TEVA BRANDED PHARM 500MG ** N012945 001

TABLET; ORAL

ACETAZOLAMIDE

AJANTA PHARMA LTD 125MG A211151 001 Sep 11, 2023

250MG A211151 002 Sep 11, 2023

ALRA 250MG A083320 001

ASCOT 250MG A087686 001 Oct 20, 1982

BRECKENRIDGE 125MG A207503 001 Apr 30, 2020

250MG A207503 002 Apr 30, 2020

HERITAGE PHARMA 250MG A088882 001 Oct 22, 1985

SUN PHARM INDUSTRIES 125MG A089753 002 Jun 22, 1988

250MG A089753 001 Jun 22, 1988

TORRENT 125MG A213706 001 Jan 26, 2024

250MG A213706 002 Jan 26, 2024

VANGARD 250MG A087654 001 Feb 05, 1982

WATSON LABS 250MG A084498 002

DIAMOX

+ TEVA BRANDED PHARM 125MG ** N008943 001

+ 250MG ** N008943 002

ACETAZOLAMIDE SODIUM

INJECTABLE; INJECTION

ACETAZOLAMIDE SODIUM

ENDO OPERATIONS EQ 500MG BASE/VIAL A205358 001 Jun 20, 2017

HOSPIRA EQ 500MG BASE/VIAL A040108 001 Oct 30, 1995

DIAMOX

+ TEVA WOMENS EQ 500MG BASE/VIAL ** N009388 001

ACETIC ACID, GLACIAL

SOLUTION/DROPS; OTIC

ACETASOL

ACTAVIS MID ATLANTIC 2% A087146 001

ACETIC ACID

CHARTWELL RX 2% A040166 001 Jul 26, 1996

KV PHARM 2% A085493 001

ORLEX

WARNER CHILCOTT 2% A086845 001

VOSOL

+ HIKMA 2% N012179 001

ACETIC ACID, GLACIAL; ALUMINUM ACETATE

SOLUTION/DROPS; OTIC

ACETIC ACID 2% IN AQUEOUS ALUMINUM ACETATE

BAUSCH AND LOMB 2%; 0.79% A040063 001 Feb 25, 1994

BOROFAIR

PHARMAFAIR 2%; 0.79% A088606 001 Aug 21, 1985

DOMEBORO

+ BAYER PHARMS 2%; 0.79% A084476 001

ACETIC ACID, GLACIAL; DESONIDE

SOLUTION/DROPS; OTIC

TRIDESILON

BAYER PHARMS 2%; 0.05% N017914 001

ACETIC ACID, GLACIAL; HYDROCORTISONE

SOLUTION/DROPS; OTIC

ACETASOL HC

ACTAVIS MID ATLANTIC 2%; 1% A087143 001 Jan 13, 1982

ACETIC ACID W/ HYDROCORTISONE

KV PHARM 2%; 1% A085492 001

HYDROCORTISONE AND ACETIC ACID

BAUSCH AND LOMB 2%; 1% A040097 001 Oct 31, 1994

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ACETIC ACID, GLACIAL; HYDROCORTISONE

SOLUTION/DROPS;OTIC

HYDROCORTISONE AND ACETIC ACID

WOCKHARDT 2%;1% A040168 001 Aug 30, 1996

ORLEX HC

WARNER CHILCOTT 2%;1% A086844 001

ACETIC ACID, GLACIAL; HYDROCORTISONE; NEOMYCIN SULFATE

SUSPENSION/DROPS;OTIC

NEO-CORT-DOME

BAYER PHARMS 2%;1%;EQ 0.35% BASE N050238 001

ACETOHEXAMIDE

TABLET;ORAL

ACETOHEXAMIDE

ANI PHARMS 250MG A070870 002 Feb 09, 1987

500MG A070870 001 Feb 09, 1987

USL PHARMA

250MG A070753 001 Nov 03, 1986

500MG A070754 001 Nov 03, 1986

WATSON LABS TEVA

250MG A071893 001 Nov 25, 1987

500MG A071894 001 Nov 25, 1987

DYMELOR

LILLY 250MG N013378 002

500MG N013378 001

ACETOPHENAZINE MALEATE

TABLET;ORAL

TINDAL

SCHERING 20MG N012254 002

ACETRIZOATE SODIUM

SOLUTION;INTRAUTERINE

SALPIX

ORTHO MCNEIL PHARM 53% N009008 001

ACETYLCOLINE CHLORIDE

FOR SOLUTION;OPHTHALMIC

MIOCHOL

+ NOVARTIS 20MG/VIAL ** N016211 001

ACETYLCYSTEINE

INJECTABLE;INTRAVENOUS

ACETYLCYSTEINE

EXELA PHARMA 6GM/30ML (200MG/ML) A204797 001 Apr 15, 2021

SOLUTION;INHALATION, ORAL

ACETYLCYSTEINE

EXELA PHARMA 20% A205643 001 Nov 01, 2023

HOSPIRA 10% A071364 001 May 01, 1989

20% A071365 001 May 01, 1989

ROXANE

10% A072323 001 Apr 30, 1992

10% A072621 001 Sep 30, 1992

20% A072324 001 Apr 30, 1992

20% A072622 001 Sep 30, 1992

MUCOMYST

+ APOTHECON 10% ** N013601 002

+ 20% ** N013601 001

MUCOSIL-10

DEY 10% A070575 001 Oct 14, 1986

MUCOSIL-20

DEY 20% A070576 001 Oct 14, 1986

TABLET, EFFERVESCENT;ORAL

CETYLEV

+ ARBOR PHARMS LLC 500MG N207916 001 Jan 29, 2016

+ 2.5GM N207916 002 Jan 29, 2016

ACETYLCYSTEINE LYSINE

FOR SOLUTION;ORAL

LEGUBETI

+ GALEPHAR EQ 500MG BASE/PACKET N215040 001 Feb 13, 2024

+ EQ 2.5GM BASE/PACKET N215040 002 Feb 13, 2024

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ACETYLCYSTEINE; ISOPROTERENOL HYDROCHLORIDE

SOLUTION; INHALATION

MUCOMYST W/ ISOPROTERENOL

MEAD JOHNSON 10%; 0.05% N017366 001

ACETYLDIGITOXIN

TABLET; ORAL

ACYLANID

NOVARTIS 0.1MG N009436 001

ACITRETIN

CAPSULE; ORAL

ACITRETIN

NORVIUM BIOSCIENCE 17.5MG A203707 001 Sep 10, 2015

22.5MG A203707 002 Sep 10, 2015

SORIATANE

+ STIEFEL LABS INC 10MG ** N019821 001 Oct 28, 1996

+ 17.5MG ** N019821 003 Aug 06, 2009

+ 22.5MG ** N019821 004 Aug 06, 2009

+ 25MG ** N019821 002 Oct 28, 1996

ACRISORCIN

CREAM; TOPICAL

AKRINOL

SCHERING 2MG/GM N012470 001

ACRIVASTINE; PSEUDOEPHEDRINE HYDROCHLORIDE

CAPSULE; ORAL

SEMPREX-D

+ ENDO OPERATIONS 8MG; 60MG N019806 001 Mar 25, 1994

ACYCLOVIR

CAPSULE; ORAL

ACYCLOVIR

ACTAVIS ELIZABETH 200MG A074906 001 Aug 26, 1997

AUROBINDO PHARMA USA 200MG A074727 001 Apr 22, 1997

CHARTWELL MOLECULES 200MG A074872 001 Apr 22, 1997

IVAX SUB TEVA PHARMS 200MG A074674 001 Apr 22, 1997

LEK PHARM 200MG A074750 001 Apr 22, 1997

MYLAN 200MG A074977 001 Apr 13, 1998

RANBAXY 200MG A074975 001 Sep 30, 1998

ROXANE 200MG A074570 002 Apr 22, 1997

STRIDES PHARMA 200MG A074833 001 Apr 22, 1997

TEVA 200MG A074828 001 Apr 22, 1997

TEVA PHARMS 200MG A074914 001 Nov 26, 1997

WATSON LABS 200MG A075101 001 Apr 15, 1998

ZOVIRAX

+ NORVIUM BIOSCIENCE 200MG ** N018828 001 Jan 25, 1985

OINTMENT; OPHTHALMIC

AVACLIR

+ FERA PHARMS LLC 3% ** N202408 001 Mar 29, 2019

OINTMENT; TOPICAL

ACYCLOVIR

ANDA REPOSITORY 5% A206437 001 Jul 31, 2017

COSETTE 5% A205591 001 Nov 13, 2017

PADAGIS ISRAEL 5% A205659 001 Feb 20, 2019

PRINSTON INC 5% A212202 001 Nov 15, 2021

SUSPENSION; ORAL

ACYCLOVIR

HIKMA 200MG/5ML A077026 001 Jun 07, 2005

VISTAPHARM LLC 200MG/5ML A213951 001 Jan 11, 2021

ZOVIRAX

+ NORVIUM BIOSCIENCE 200MG/5ML N019909 001 Dec 22, 1989

TABLET; ORAL

ACYCLOVIR

ACTAVIS ELIZABETH 400MG A074870 001 Jun 05, 1997

800MG A074870 002 Jun 05, 1997

AUROBINDO PHARMA USA 400MG A075211 001 Sep 28, 1998

800MG A075211 002 Sep 28, 1998

CHARTWELL MOLECULES 400MG A074834 001 Apr 24, 1997

800MG A074834 002 Apr 24, 1997

IVAX SUB TEVA PHARMS 400MG A074836 001 Apr 22, 1997

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ACYCLOVIR

TABLET; ORAL

ACYCLOVIR

	800MG	A074836 002	Apr 22, 1997
LEK PHARM	400MG	A074658 001	Apr 22, 1997
	800MG	A074658 002	Apr 22, 1997
MYLAN	400MG	A074976 001	Apr 13, 1998
	800MG	A074976 002	Apr 13, 1998
SUN PHARM INDS LTD	400MG	A074980 001	Sep 30, 1998
	800MG	A074980 002	Sep 30, 1998
TEVA	200MG **	A074556 001	Apr 22, 1997
TEVA PHARMS	400MG	A075021 001	Mar 18, 1998
	800MG	A075021 002	Mar 18, 1998
ZOVIRAX			
+ NORVIUM BIOSCIENCE	400MG **	N020089 001	Apr 30, 1991
+	800MG **	N020089 002	Apr 30, 1991

ACYCLOVIR SODIUM

INJECTABLE; INJECTION

ACYCLOVIR

ABBVIE	EQ 50MG BASE/ML	A075114 001	Jul 26, 1999
ACYCLOVIR IN SODIUM CHLORIDE 0.9% PRESERVATIVE FREE			
EUROHLTH INTL SARL	EQ 500MG BASE/VIAL	A074885 001	Dec 19, 1997
	EQ 1GM BASE/VIAL	A074885 002	Dec 19, 1997
ACYCLOVIR SODIUM			
APOTHECON	EQ 500MG BASE/VIAL	A074897 001	Feb 27, 1998
	EQ 1GM BASE/VIAL	A074897 002	Feb 27, 1998
CHARTWELL INJECTABLE	EQ 500MG BASE/VIAL	A074596 002	Apr 22, 1997
	EQ 1GM BASE/VIAL	A074596 001	Apr 22, 1997
DR REDDYS	EQ 50MG BASE/ML	A207919 001	Jun 17, 2020
EUROHLTH INTL SARL	EQ 500MG BASE/VIAL	A074913 001	Oct 15, 1997
	EQ 1GM BASE/VIAL	A074913 002	Oct 15, 1997
FRESENIUS KABI USA	EQ 500MG BASE/VIAL	A075015 001	Apr 30, 1998
HIKMA	EQ 500MG BASE/VIAL	A205771 001	Feb 29, 2016
	EQ 1GM BASE/VIAL	A205771 002	Feb 29, 2016
HOSPIRA	EQ 25MG BASE/ML	A074720 001	Apr 22, 1997
	EQ 50MG BASE/ML	A075065 001	Feb 25, 1999
	EQ 500MG BASE/VIAL	A074663 001	Apr 22, 1997
	EQ 500MG BASE/VIAL	A074758 001	Apr 22, 1997
	EQ 1GM BASE/VIAL	A074663 002	Apr 22, 1997
	EQ 1GM BASE/VIAL	A074758 002	Apr 22, 1997
NORVIUM BIOSCIENCE	EQ 500MG BASE/VIAL	A203927 001	Mar 29, 2017
	EQ 1GM BASE/VIAL	A203927 002	Mar 29, 2017
TEVA PARENTERAL	EQ 50MG BASE/ML	A075627 001	Mar 28, 2001
	EQ 500MG BASE/VIAL	A074969 001	Aug 26, 1997
	EQ 1GM BASE/VIAL	A074969 002	Aug 26, 1997
ZYDUS PHARMS	EQ 500MG BASE/VIAL	A206606 001	Jun 13, 2017
	EQ 1GM BASE/VIAL	A206606 002	Jun 13, 2017
ZOVIRAX			
+ GLAXOSMITHKLINE	EQ 250MG BASE/VIAL **	N018603 003	Aug 30, 1983
+	EQ 500MG BASE/VIAL **	N018603 001	Oct 22, 1982
+	EQ 1GM BASE/VIAL **	N018603 002	Jun 29, 1989

ADAPALENE

LOTION; TOPICAL

DIFFERIN

+ GALDERMA LABS LP	0.1%	N022502 001	Mar 17, 2010
--------------------	------	-------------	--------------

SOLUTION; TOPICAL

DIFFERIN

+ GALDERMA LABS LP	0.1% **	N020338 001	May 31, 1996
--------------------	---------	-------------	--------------

ADAPALENE; BENZOYL PEROXIDE

GEL; TOPICAL

ADAPALENE AND BENZOYL PEROXIDE

ACTAVIS MID ATLANTIC	0.1%; 2.5%	A203790 001	Sep 30, 2015
----------------------	------------	-------------	--------------

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ADEFOVIR DIPIVOXIL

TABLET; ORAL

HEPSERA

+ GILEAD 10MG ** N021449 001 Sep 20, 2002

ADENOSINE

INJECTABLE; INJECTION

ADENOCARD

+ ASTELLAS 3MG/ML ** N019937 002 Oct 30, 1989

ADENOSINE

AM REGENT 3MG/ML A090010 001 Apr 28, 2009

HIKMA 3MG/ML A076501 001 Jun 16, 2004

NORVIUM BIOSCIENCE 3MG/ML A078640 001 Mar 21, 2014

TEVA PHARMS USA 3MG/ML A076564 001 Jun 16, 2004

WOCKHARDT 3MG/ML A078676 001 Jul 31, 2008

WOCKHARDT 3MG/ML A090220 001 Jul 20, 2009

SOLUTION; INTRAVENOUS

ADENOSCAN

+ ASTELLAS 60MG/20ML (3MG/ML) ** N020059 001 May 18, 1995

+ 90MG/30ML (3MG/ML) ** N020059 002 May 18, 1995

ADENOSINE

HOSPIRA 60MG/20ML (3MG/ML) A203883 001 Mar 24, 2014

90MG/30ML (3MG/ML) A203883 002 Mar 24, 2014

ALATROFLOXACIN MESYLATE

INJECTABLE; INJECTION

TROVAN PRESERVATIVE FREE

PFIZER EQ 200MG BASE/VIAL N020760 001 Dec 18, 1997

EQ 300MG BASE/VIAL N020760 002 Dec 18, 1997

ALBENDAZOLE

TABLET; ORAL

ALBENDAZOLE

CHARTWELL RX 200MG A211636 001 Jun 10, 2020

CIPLA LTD 200MG A210434 001 Sep 21, 2018

STRIDES PHARMA 200MG A210011 001 Dec 07, 2018

ALBENZA

+ IMPAX LABS INC 200MG ** N020666 001 Jun 11, 1996

ALBUTEROL

AEROSOL, METERED; INHALATION

ALBUTEROL

ARMSTRONG PHARMS 0.09MG/INH A072273 001 Aug 14, 1996

GENPHARM 0.09MG/INH A073045 001 Aug 19, 1997

IVAX SUB TEVA PHARMS 0.09MG/INH A073272 001 Dec 28, 1995

PLIVA 0.09MG/INH A074072 001 Aug 01, 1996

PROVENTIL

SCHERING 0.09MG/INH N017559 001

VENTOLIN

GLAXOSMITHKLINE 0.09MG/INH N018473 001

ALBUTEROL SULFATE

AEROSOL, METERED; INHALATION

ALBUTEROL SULFATE

PADAGIS US EQ 0.09MG BASE/INH A203760 001 Feb 24, 2020

CAPSULE; INHALATION

VENTOLIN ROTACAPS

GLAXOSMITHKLINE EQ 0.2MG BASE N019489 001 May 04, 1988

POWDER, METERED; INHALATION

PROAIR DIGIHALER

+ TEVA BRANDED PHARM EQ 0.09MG BASE/INH N205636 002 Dec 21, 2018

SOLUTION; INHALATION

ACCUNEB

+ NORVIUM BIOSCIENCE EQ 0.021% BASE ** N020949 002 Apr 30, 2001

+ EQ 0.042% BASE ** N020949 001 Apr 30, 2001

ALBUTEROL SULFATE

ACTAVIS MID ATLANTIC EQ 0.083% BASE A073533 001 Sep 26, 1995

APOTEX INC EQ 0.021% BASE A078623 001 Apr 05, 2010

EQ 0.042% BASE A078623 002 Apr 05, 2010

EQ 0.083% BASE A075717 001 Feb 02, 2007

EQ 0.5% BASE A076391 001 Apr 01, 2003

BAUSCH EQ 0.083% BASE A075358 001 Mar 29, 2000

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ALBUTEROL SULFATE

SOLUTION; INHALATION

ALBUTEROL SULFATE

		EQ 0.5% BASE	A075050	001	Jun 18, 1998
	COPLEY PHARM	EQ 0.083% BASE	A073495	001	May 28, 1993
		EQ 0.5% BASE	A073307	001	Nov 27, 1991
	EPIC PHARMA LLC	EQ 0.083% BASE	A075063	001	Feb 09, 1999
	LANDELA PHARM	EQ 0.083% BASE	A077569	001	Apr 04, 2006
	NORVIUM BIOSCIENCE	EQ 0.083% BASE **	A072652	001	Feb 21, 1992
	ROXANE	EQ 0.083% BASE	A075129	001	Feb 13, 2001
	TEVA PHARMS	EQ 0.083% BASE	A075343	001	Nov 09, 1999
	WATSON LABS	EQ 0.021% BASE	A077772	001	Sep 25, 2007
		EQ 0.042% BASE	A077772	002	Sep 25, 2007
	WATSON LABS INC	EQ 0.083% BASE	A076370	001	Nov 24, 2003
	WOCKHARDT EU OPERATN	EQ 0.083% BASE	A075394	001	Nov 22, 1999
	PROVENTIL				
	+ SCHERING	EQ 0.083% BASE **	N019243	002	Jan 14, 1987
	+	EQ 0.5% BASE **	N019243	001	Jan 14, 1987
	VENTOLIN				
	+ GLAXOSMITHKLINE	EQ 0.083% BASE **	N019773	001	Apr 23, 1992
		EQ 0.5% BASE **	N019269	002	Jan 16, 1987
	SYRUP; ORAL				
	ALBUTEROL SULFATE				
	ACTAVIS MID ATLANTIC	EQ 2MG BASE/5ML	A075262	001	Mar 30, 1999
	HIKMA	EQ 2MG BASE/5ML	A074749	001	Jan 30, 1998
	MOVA	EQ 2MG BASE/5ML	A074302	001	Sep 30, 1994
	WATSON LABS	EQ 2MG BASE/5ML	A073165	001	Apr 29, 1993
	PROVENTIL				
	+ SCHERING	EQ 2MG BASE/5ML **	N018062	001	Jan 19, 1983
	VENTOLIN				
	GLAXOSMITHKLINE	EQ 2MG BASE/5ML **	N019621	001	Jun 10, 1987
	TABLET; ORAL				
	ALBUTEROL SULFATE				
	AM THERAP	EQ 2MG BASE	A072449	001	Dec 05, 1989
		EQ 4MG BASE	A072450	001	Dec 05, 1989
	AUROBINDO PHARMA LTD	EQ 2MG BASE	A213657	001	May 14, 2020
		EQ 4MG BASE	A213657	002	May 14, 2020
	CHARTWELL RX	EQ 2MG BASE	A072151	001	Dec 05, 1989
		EQ 4MG BASE	A072151	002	Dec 05, 1989
	COPLEY PHARM	EQ 2MG BASE	A072966	001	Nov 22, 1991
		EQ 4MG BASE	A072967	001	Nov 22, 1991
	HIBROW HLTHCARE	EQ 2MG BASE	A213524	001	Oct 08, 2020
		EQ 4MG BASE	A213524	002	Oct 08, 2020
	PLIVA	EQ 2MG BASE	A072316	001	Dec 05, 1989
		EQ 4MG BASE	A072317	001	Dec 05, 1989
	STRIDES PHARMA	EQ 2MG BASE	A072860	002	Dec 20, 1989
		EQ 4MG BASE	A072860	001	Dec 20, 1989
	TEVA	EQ 2MG BASE	A072619	001	Dec 05, 1989
		EQ 2MG BASE	A072779	001	Jun 25, 1993
		EQ 2MG BASE	A072938	001	Mar 30, 1990
		EQ 4MG BASE	A072620	001	Dec 05, 1989
		EQ 4MG BASE	A072780	001	Jun 25, 1993
		EQ 4MG BASE	A072939	001	Mar 30, 1990
	UCB INC	EQ 2MG BASE	A073120	001	Sep 29, 1992
		EQ 4MG BASE	A073121	001	Sep 29, 1992
	WARNER CHILCOTT	EQ 2MG BASE	A072817	001	Jan 09, 1990
		EQ 4MG BASE	A072818	001	Jan 09, 1990
	WATSON LABS	EQ 2MG BASE	A072629	001	Jan 31, 1991
		EQ 2MG BASE	A072764	001	Aug 28, 1991
		EQ 4MG BASE	A072630	001	Jan 31, 1991
		EQ 4MG BASE	A072765	001	Aug 28, 1991
	PROVENTIL				
	+ SCHERING	EQ 2MG BASE **	N017853	001	May 07, 1982
	+	EQ 4MG BASE **	N017853	002	May 07, 1982
	VENTOLIN				
	GLAXOSMITHKLINE	EQ 2MG BASE	N019112	001	Jul 10, 1986
		EQ 4MG BASE	N019112	002	Jul 10, 1986

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ALBUTEROL SULFATE

TABLET, EXTENDED RELEASE;ORAL

ALBUTEROL SULFATE

RISING	EQ 4MG BASE	A078092 002	Jan 29, 2007
	EQ 8MG BASE	A078092 001	Jan 29, 2007
PROVENTIL			
SCHERING	EQ 4MG BASE	N019383 001	Jul 13, 1987
VOLMAX			
+ MURO	EQ 4MG BASE	N019604 002	Dec 23, 1992
+	EQ 8MG BASE	N019604 001	Dec 23, 1992
VOSPIRE ER			
STRIDES PHARMA	EQ 4MG BASE	A076130 002	Sep 26, 2002
	EQ 8MG BASE	A076130 003	Sep 26, 2002

ALBUTEROL SULFATE; IPRATROPIUM BROMIDE

AEROSOL, METERED; INHALATION

COMBIVENT

BOEHRINGER INGELHEIM	EQ 0.09MG BASE/INH;0.018MG/INH	N020291 001	Oct 24, 1996
SOLUTION; INHALATION			
ALBUTEROL SULFATE AND IPRATROPIUM BROMIDE			
AIPING PHARM INC	EQ 0.083% BASE;0.017%	A076867 001	Dec 21, 2006
APOTEX INC	EQ 0.083% BASE;0.017%	A077117 001	Dec 31, 2007
LUOXIN AUROVITAS	EQ 0.083% BASE;0.017%	A206532 001	Jul 08, 2020
TEVA PHARMS	EQ 0.083% BASE;0.017%	A076724 001	Dec 31, 2007
WATSON LABS TEVA	EQ 0.083% BASE;0.017%	A077063 001	Dec 31, 2007
DUONEB			
+ NORVIUM BIOSCIENCE	EQ 0.083% BASE;0.017% **	N020950 001	Mar 21, 2001

ALCAFTADINE

SOLUTION/DROPS;OPHTHALMIC

ALCAFTADINE

EUGIA PHARMA	0.25%	A210659 001	Jun 23, 2023
--------------	-------	-------------	--------------

ALCLOMETASONE DIPROPIONATE

CREAM; TOPICAL

ACLOVATE

+ FOUGERA PHARMS	0.05% **	N018707 001	Dec 14, 1982
------------------	----------	-------------	--------------

OINTMENT; TOPICAL

ACLOVATE

+ FOUGERA PHARMS	0.05% **	N018702 001	Dec 14, 1982
------------------	----------	-------------	--------------

ALCOHOL

SOLUTION; INTRA-ARTERIAL

ABLYSINOL

+ BPI LABS	99% (1ML)	N207987 001	Jun 21, 2018
------------	-----------	-------------	--------------

ALENDRONATE SODIUM

SOLUTION; ORAL

FOSAMAX

+ MERCK	EQ 70MG BASE/75ML **	N021575 001	Sep 17, 2003
---------	----------------------	-------------	--------------

TABLET; ORAL

ALENDRONATE SODIUM

CHARTWELL RX	EQ 5MG BASE	A075871 001	Apr 22, 2009
	EQ 5MG BASE	A079049 003	Aug 04, 2008
	EQ 10MG BASE	A075871 002	Apr 22, 2009
	EQ 10MG BASE	A079049 004	Aug 04, 2008
	EQ 35MG BASE	A075871 004	Apr 22, 2009
	EQ 35MG BASE	A079049 001	Aug 04, 2008
	EQ 40MG BASE	A075871 003	Apr 22, 2009
	EQ 70MG BASE	A075871 005	Apr 22, 2009
	EQ 70MG BASE	A079049 002	Aug 04, 2008
IMPAX LABS INC	EQ 5MG BASE	A075710 001	Feb 06, 2008
	EQ 10MG BASE	A075710 002	Feb 06, 2008
	EQ 35MG BASE	A075710 003	Feb 06, 2008
	EQ 40MG BASE	A075710 004	Feb 06, 2008
	EQ 70MG BASE	A075710 005	Feb 06, 2008
JUBILANT CADISTA	EQ 5MG BASE	A090557 001	Feb 18, 2010
	EQ 10MG BASE	A090557 002	Feb 18, 2010
	EQ 35MG BASE	A090557 003	Feb 18, 2010
	EQ 70MG BASE	A090557 004	Feb 18, 2010
MYLAN	EQ 35MG BASE	A078638 001	Aug 04, 2008
	EQ 70MG BASE	A078638 002	Aug 04, 2008

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ALENDRONATE SODIUM

TABLET;ORAL

ALENDRONATE SODIUM

RISING	EQ 5MG BASE	A076584 001	Aug 04, 2008
	EQ 10MG BASE	A076584 002	Aug 04, 2008
	EQ 35MG BASE	A076584 003	Aug 04, 2008
	EQ 70MG BASE	A076584 004	Aug 04, 2008
TEVA PHARMS	EQ 35MG BASE	A076184 002	Aug 04, 2008
	EQ 70MG BASE	A076184 001	Feb 06, 2008

FOSAMAX

+ ORGANON	EQ 5MG BASE **	N020560 003	Apr 25, 1997
+	EQ 10MG BASE **	N020560 001	Sep 29, 1995
+	EQ 35MG BASE **	N020560 004	Oct 20, 2000
+	EQ 40MG BASE **	N020560 002	Sep 29, 1995

ALFUZOSIN HYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL

ALFUZOSIN HYDROCHLORIDE

CHARTWELL RX	10MG	A079056 001	Jul 18, 2011
NORVIUM BIOSCIENCE	10MG	A079014 001	Jul 18, 2011
SUN PHARM	10MG	A079057 001	Jul 18, 2011
TORRENT PHARMS	10MG	A079054 001	Jul 18, 2011
WOCKHARDT	10MG	A090221 001	Aug 10, 2012

ALISKIREN HEMIFUMARATE

CAPSULE, PELLET;ORAL

TEKTURNA

+ NODEN PHARMA	EQ 37.5MG BASE	N210709 001	Nov 14, 2017
----------------	----------------	-------------	--------------

ALISKIREN HEMIFUMARATE; AMLODIPINE BESYLATE

TABLET;ORAL

TEKAMLO

NOVARTIS	EQ 150MG BASE;EQ 5MG BASE	N022545 001	Aug 26, 2010
	EQ 150MG BASE;EQ 10MG BASE	N022545 002	Aug 26, 2010
	EQ 300MG BASE;EQ 5MG BASE	N022545 003	Aug 26, 2010
	EQ 300MG BASE;EQ 10MG BASE	N022545 004	Aug 26, 2010

ALISKIREN HEMIFUMARATE; AMLODIPINE BESYLATE; HYDROCHLOROTHIAZIDE

TABLET;ORAL

AMTURNIDE

NOVARTIS	EQ 150MG BASE;EQ 5MG BASE;12.5MG	N200045 001	Dec 21, 2010
	EQ 300MG BASE;EQ 5MG BASE;12.5MG	N200045 002	Dec 21, 2010
	EQ 300MG BASE;EQ 5MG BASE;25MG	N200045 003	Dec 21, 2010
	EQ 300MG BASE;EQ 10MG BASE;12.5MG	N200045 004	Dec 21, 2010
	EQ 300MG BASE;EQ 10MG BASE;25MG	N200045 005	Dec 21, 2010

ALISKIREN HEMIFUMARATE; HYDROCHLOROTHIAZIDE

TABLET;ORAL

TEKTURNA HCT

+ NODEN PHARMA	EQ 150MG BASE;12.5MG **	N022107 001	Jan 18, 2008
+	EQ 150MG BASE;25MG **	N022107 002	Jan 18, 2008
+	EQ 300MG BASE;12.5MG **	N022107 003	Jan 18, 2008
+	EQ 300MG BASE;25MG **	N022107 004	Jan 18, 2008

ALISKIREN HEMIFUMARATE; VALSARTAN

TABLET;ORAL

VALTURNA

NOVARTIS	EQ 150MG BASE;160MG	N022217 001	Sep 16, 2009
	EQ 300MG BASE;320MG	N022217 002	Sep 16, 2009

ALKAVERVIR

TABLET;ORAL

VERILOID

3M	2MG	N007336 002
	3MG	N007336 003

ALLOPURINOL

TABLET;ORAL

ALLOPURINOL

AIPING PHARM INC	100MG	A070268 001	Dec 31, 1985
IPCA LABS LTD	100MG	A090637 001	Mar 16, 2011
	300MG	A090637 002	Mar 16, 2011
LUPIN LTD	100MG	A211807 001	Dec 14, 2023

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ALLOPURINOL

TABLET;ORAL

ALLOPURINOL

	300MG	A211807 002	Dec 14, 2023
MUTUAL PHARM	100MG	A070466 001	Dec 24, 1985
	300MG	A070467 001	Dec 24, 1985
PURACAP PHARM	100MG	A070150 001	Dec 10, 1985
	300MG	A070147 001	Dec 10, 1985
PUREPAC PHARM	100MG	A070579 001	Apr 14, 1986
	300MG	A070580 001	Apr 14, 1986
SANDOZ	300MG	A070269 001	Dec 31, 1985
SUN PHARM INDS INC	100MG	A078390 001	Aug 30, 2007
	300MG	A078390 002	Aug 30, 2007
SUPERPHARM	100MG	A070950 001	Nov 30, 1988
	300MG	A070951 001	Nov 30, 1988
WATSON LABS	100MG	N018241 001	Nov 16, 1984
	100MG	N018785 001	Sep 28, 1984
	300MG	N018241 002	Nov 16, 1984
	300MG	N018785 002	Sep 28, 1984

LOPURIN

ABBOTT

100MG	N018297 001
300MG	N018297 002

ALLOPURINOL; LESINURAD

TABLET;ORAL

DUZALLO

+ IRONWOOD PHARMS INC	200MG;200MG	N209203 001	Aug 18, 2017
+	300MG;200MG	N209203 002	Aug 18, 2017

ALMOTRIPTAN MALATE

TABLET;ORAL

AXERT

+ JANSSEN PHARMS	EQ 6.25MG BASE **	N021001 001	May 07, 2001
+	EQ 12.5MG BASE **	N021001 002	May 07, 2001

ALOGLIPTIN BENZOATE; PIOGLITAZONE HYDROCHLORIDE

TABLET;ORAL

OSEN

+ TAKEDA PHARMS USA	EQ 12.5MG BASE;EQ 15MG BASE **	N022426 004	Jan 25, 2013
+	EQ 12.5MG BASE;EQ 45MG BASE **	N022426 006	Jan 25, 2013

ALOSETRON HYDROCHLORIDE

TABLET;ORAL

ALOSETRON HYDROCHLORIDE

ENDO OPERATIONS	EQ 0.5MG BASE	A206113 001	Feb 23, 2018
	EQ 1MG BASE	A206113 002	Feb 23, 2018
HIBROW HLTHCARE	EQ 0.5MG BASE	A211621 001	Sep 16, 2019
	EQ 1MG BASE	A211621 002	Sep 16, 2019
HIKMA	EQ 0.5MG BASE	A200652 001	May 04, 2015
	EQ 1MG BASE	A200652 002	May 04, 2015

ALPRAZOLAM

SOLUTION;ORAL

ALPRAZOLAM

ROXANE	0.5MG/5ML	A074314 001	Oct 31, 1993
--------	-----------	-------------	--------------

TABLET;ORAL

ALPRAZOLAM

ANI PHARMS	0.25MG	A074085 001	Feb 16, 1994
	0.5MG	A074085 002	Feb 16, 1994
	1MG	A074085 003	Feb 16, 1994
	2MG	A074085 004	Feb 26, 1996
IVAX SUB TEVA PHARMS	0.25MG	A074294 001	Jul 29, 1994
	0.5MG	A074294 002	Jul 29, 1994
	1MG	A074294 003	Jul 29, 1994
	2MG	A074294 004	Jul 29, 1994
MYLAN	0.25MG	A074215 001	Jan 27, 1994
	0.5MG	A074215 002	Jan 27, 1994
	1MG	A074215 003	Jan 27, 1994
	2MG	A074215 004	Jan 27, 1994
NORVIUM BIOSCIENCE	0.25MG	A074046 001	Oct 19, 1993
	0.5MG	A074046 002	Oct 19, 1993
	1MG	A074046 003	Oct 19, 1993

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ALPRAZOLAMTABLET; ORAL
ALPRAZOLAM

	2MG	A074046 004	May 07, 1997
OXFORD PHARMS	0.25MG	A078491 001	Sep 25, 2008
	0.5MG	A078491 002	Sep 25, 2008
	1MG	A078491 003	Sep 25, 2008
	2MG	A078491 004	Dec 12, 2008
ROXANE	0.25MG	A074199 001	Oct 19, 1993
	0.5MG	A074199 002	Oct 19, 1993
	1MG	A074199 003	Oct 19, 1993
WATSON LABS	0.25MG	A074456 001	Aug 31, 1995
	0.25MG	A074479 001	Jan 21, 1997
	0.5MG	A074456 002	Aug 31, 1995
	0.5MG	A074479 002	Jan 21, 1997
	1MG	A074456 003	Aug 31, 1995
	1MG	A074479 003	Jan 21, 1997

TABLET, EXTENDED RELEASE; ORAL
ALPRAZOLAM

ACTAVIS LABS FL INC	0.5MG	A077198 001	May 13, 2010
	1MG	A077198 002	May 13, 2010
	2MG	A077198 003	May 13, 2010
	3MG	A077198 004	May 13, 2010
ANI PHARMS	0.5MG	A077725 001	Jul 31, 2006
	0.5MG	A077979 001	Feb 28, 2007
	1MG	A077725 002	Jul 31, 2006
	1MG	A077979 002	Feb 28, 2007
	2MG	A077725 004	Jul 31, 2006
	2MG	A077979 003	Feb 28, 2007
	3MG	A077725 003	Jul 31, 2006
	3MG	A077979 004	Feb 28, 2007
ENDO OPERATIONS	0.5MG	A078442 001	Oct 15, 2007
	0.5MG	A078469 001	Sep 29, 2011
	1MG	A078442 002	Oct 15, 2007
	1MG	A078469 002	Sep 29, 2011
	2MG	A078442 003	Oct 15, 2007
	2MG	A078469 003	Sep 29, 2011
	3MG	A078442 004	Oct 15, 2007
	3MG	A078469 004	Sep 29, 2011
HERITAGE PHARMS INC	0.5MG	A078489 001	Oct 17, 2008
	1MG	A078489 002	Oct 17, 2008
	2MG	A078489 003	Oct 17, 2008
	3MG	A078489 004	Oct 17, 2008
IMPAX LABS	0.5MG	A077968 004	May 24, 2007
	1MG	A077968 003	May 24, 2007
	2MG	A077968 002	May 24, 2007
	3MG	A077968 001	May 24, 2007
IMPAX LABS INC	0.5MG	A077996 001	Jan 31, 2007
	1MG	A077996 002	Jan 31, 2007
	2MG	A077996 003	Jan 31, 2007
	3MG	A077996 004	Jan 31, 2007
NORVIUM BIOSCIENCE	0.5MG	A077391 002	Jan 26, 2006
	1MG	A077391 003	Jan 26, 2006
	2MG	A077391 004	Jan 26, 2006
	3MG	A077391 001	Jan 26, 2006
SANDOZ INC	0.5MG	A077777 001	Jun 30, 2006
	1MG	A077777 002	Jun 30, 2006
	2MG	A077777 003	Jun 30, 2006
	3MG	A077777 004	Jun 30, 2006

TABLET, ORALLY DISINTEGRATING; ORAL
NIRAVAM

+	UCB INC	0.25MG **	N021726 001	Jan 19, 2005
+		0.5MG **	N021726 002	Jan 19, 2005
+		1MG **	N021726 003	Jan 19, 2005
+		2MG **	N021726 004	Jan 19, 2005

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ALPROSTADIL

INJECTABLE; INJECTION

CAVERJECT

+	PFIZER	0.005MG/VIAL	N020379	003	Jun 27, 1996
		0.005MG/ML	N020755	001	Oct 31, 1997
		0.01MG/ML	N020755	002	Oct 01, 1997
		0.02MG/ML	N020755	003	Oct 01, 1997

EDEX

	ENDO OPERATIONS	0.005MG/VIAL	N020649	001	Jun 12, 1997
--	-----------------	--------------	---------	-----	--------------

SUPPOSITORY; URETHRAL

MUSE

+	MYLAN SPECIALITY LP	0.125MG	N020700	001	Nov 19, 1996
+		0.25MG	N020700	002	Nov 19, 1996
+		0.5MG	N020700	003	Nov 19, 1996
+		1MG	N020700	004	Nov 19, 1996

ALSEROXYLON

TABLET; ORAL

RAUTENSIN

	NOVARTIS	2MG	N009215	001	
--	----------	-----	---------	-----	--

RAUWILOID

	3M	2MG	N008867	001	
--	----	-----	---------	-----	--

ALTRETAMINE

CAPSULE; ORAL

HEXALEN

+	EISAI INC	50MG	N019926	001	Dec 26, 1990
---	-----------	------	---------	-----	--------------

ALUMINUM HYDROXIDE; MAGNESIUM TRISILICATE

TABLET, CHEWABLE; ORAL

ALUMINUM HYDROXIDE AND MAGNESIUM TRISILICATE

	PENNEX	80MG; 20MG	A089449	001	Nov 27, 1987
--	--------	------------	---------	-----	--------------

FOAMCOAT

	GUARDIAN DRUG	80MG; 20MG	A071793	001	Sep 04, 1987
--	---------------	------------	---------	-----	--------------

FOAMICON

	NOVARTIS	80MG; 20MG	A072687	001	Jun 28, 1989
--	----------	------------	---------	-----	--------------

GAVISCON

+	CHATTEM SANOFI	80MG; 20MG **	N018685	001	Dec 09, 1983
---	----------------	---------------	---------	-----	--------------

+		160MG; 40MG **	N018685	002	Dec 09, 1983
---	--	----------------	---------	-----	--------------

ALVIMOPAN

CAPSULE; ORAL

ENTEREG

+	CUBIST PHARMS	12MG **	N021775	001	May 20, 2008
---	---------------	---------	---------	-----	--------------

AMANTADINE HYDROCHLORIDE

CAPSULE; ORAL

AMANTADINE HYDROCHLORIDE

	ACTAVIS ELIZABETH	100MG	A077659	001	Feb 23, 2006
--	-------------------	-------	---------	-----	--------------

	CHARTWELL MOLECULAR	100MG	A209221	001	Jun 15, 2017
--	---------------------	-------	---------	-----	--------------

	INVAGEN PHARMS	100MG	A207570	001	Sep 30, 2016
--	----------------	-------	---------	-----	--------------

	WATSON LABS	100MG	A071382	001	Jan 21, 1987
--	-------------	-------	---------	-----	--------------

	WATSON LABS INC	100MG	A208107	001	Dec 06, 2016
--	-----------------	-------	---------	-----	--------------

SYMADINE

	SOLVAY	100MG	A071000	001	Sep 04, 1986
--	--------	-------	---------	-----	--------------

SYMMETREL

+	ENDO PHARMS	100MG **	N016020	001	
---	-------------	----------	---------	-----	--

SYRUP; ORAL

AMANTADINE HYDROCHLORIDE

	CHARTWELL RX	50MG/5ML	A076352	001	Sep 10, 2004
--	--------------	----------	---------	-----	--------------

	CMP PHARMA INC	50MG/5ML	A075819	001	Sep 11, 2002
--	----------------	----------	---------	-----	--------------

	ENDO OPERATIONS	50MG/5ML	A077992	001	Dec 12, 2006
--	-----------------	----------	---------	-----	--------------

	G AND W LABS INC	50MG/5ML	A072655	001	Oct 30, 1990
--	------------------	----------	---------	-----	--------------

	TEVA PHARMS	50MG/5ML	A073115	001	Aug 23, 1991
--	-------------	----------	---------	-----	--------------

SYMMETREL

+	ENDO PHARMS	50MG/5ML **	N016023	002	
---	-------------	-------------	---------	-----	--

TABLET; ORAL

AMANTADINE HYDROCHLORIDE

	ADAPTIS	100MG	A212407	001	May 27, 2022
--	---------	-------	---------	-----	--------------

	INVAGEN PHARMS	100MG	A207571	001	Jan 31, 2017
--	----------------	-------	---------	-----	--------------

	JUBILANT GENERICS	100MG	A210403	001	Feb 07, 2018
--	-------------------	-------	---------	-----	--------------

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

AMANTADINE HYDROCHLORIDE

TABLET; ORAL

SYMMETREL

+ ENDO PHARMS 100MG ** N018101 001

TABLET, EXTENDED RELEASE; ORAL

OSMOLEX ER

+ SUPERNUS PHARMS EQ 129MG BASE N209410 001 Feb 16, 2018

+ EQ 161MG BASE N209410 004 Apr 22, 2020

+ EQ 193MG BASE N209410 002 Feb 16, 2018

+ EQ 258MG BASE N209410 003 Feb 16, 2018

AMBENONIUM CHLORIDE

TABLET; ORAL

MYTELASE

SANOFI AVENTIS US 10MG N010155 002

AMBRISENTAN

TABLET; ORAL

AMBRISENTAN

ENDO OPERATIONS 5MG A209509 001 Apr 10, 2019

10MG A209509 002 Apr 10, 2019

AMCINONIDE

CREAM; TOPICAL

CYCLOCORT

+ ASTELLAS 0.025% ** N018116 001

+ 0.1% ** N018116 002

LOTION; TOPICAL

CYCLOCORT

+ ASTELLAS 0.1% N019729 001 Jun 13, 1988

OINTMENT; TOPICAL

CYCLOCORT

+ ASTELLAS 0.1% ** N018498 001

AMDINOCILLIN

INJECTABLE; INJECTION

COACTIN

ROCHE 250MG/VIAL N050565 001 Dec 21, 1984

500MG/VIAL N050565 002 Dec 21, 1984

1GM/VIAL N050565 003 Dec 21, 1984

AMIFOSTINE

INJECTABLE; INJECTION

AMIFOSTINE

EUGIA PHARMA SPECLTS 500MG/VIAL A204363 001 Jul 17, 2017

SUN PHARM 500MG/VIAL A077126 001 Mar 14, 2008

ETHYOL

COSETTE 375MG/VIAL N020221 002 Sep 10, 1999

+ 500MG/VIAL N020221 001 Dec 08, 1995

AMIKACIN SULFATE

INJECTABLE; INJECTION

AMIKACIN SULFATE

ABBOTT EQ 250MG BASE/ML A063265 001 Nov 30, 1994

EQ 250MG BASE/ML A063266 001 Oct 31, 1994

FRESENIUS KABI USA EQ 50MG BASE/ML A205605 001 Dec 09, 2015

HIKMA EQ 50MG BASE/ML A063274 001 May 18, 1992

EQ 250MG BASE/ML A063275 001 May 18, 1992

HOSPIRA EQ 50MG BASE/ML A063263 001 Nov 30, 1994

EQ 50MG BASE/ML A063350 001 Jul 30, 1993

EQ 62.5MG BASE/ML A063283 001 Oct 31, 1994

EQ 250MG BASE/ML A063264 001 Nov 30, 1994

EQ 250MG BASE/ML A063350 002 Jul 30, 1993

EQ 250MG BASE/ML A064098 001 Jun 26, 1995

EQ 250MG BASE/ML A064099 001 Jun 20, 1995

IGI LABS INC EQ 50MG BASE/ML A063167 001 Dec 14, 1995

EQ 250MG BASE/ML A063169 001 Dec 14, 1995

MEITHEAL EQ 50MG BASE/ML A064045 001 Sep 28, 1993

AMIKACIN SULFATE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

HOSPIRA EQ 500MG BASE/100ML A064146 001 Apr 02, 1997

AMIKIN

APOTHECON EQ 50MG BASE/ML A062311 001

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

AMIKACIN SULFATE

INJECTABLE; INJECTION

AMIKIN

	EQ 50MG BASE/ML	A062562 001	Sep 20, 1984
+	EQ 50MG BASE/ML **	N050495 001	
	EQ 250MG BASE/ML	A062311 002	
	EQ 250MG BASE/ML	A062562 002	Sep 20, 1984
+	EQ 250MG BASE/ML **	N050495 002	
AMIKIN IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER			
APOTHECON			
	EQ 5MG BASE/ML	N050618 002	Nov 30, 1987
	EQ 10MG BASE/ML	N050618 001	Nov 30, 1987

AMILORIDE HYDROCHLORIDE

TABLET; ORAL

AMILORIDE HYDROCHLORIDE

CHARTWELL RX	5MG	A204180 001	Aug 07, 2015
--------------	-----	-------------	--------------

AMILORIDE HYDROCHLORIDE; HYDROCHLOROTHIAZIDE

TABLET; ORAL

AMILORIDE HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE

CHARTWELL RX	EQ 5MG ANHYDROUS; 50MG	A073357 001	Nov 27, 1991
TEVA	EQ 5MG ANHYDROUS; 50MG	A070795 001	Apr 17, 1988
WATSON LABS	EQ 5MG ANHYDROUS; 50MG	A073334 001	Jul 19, 1991
HYDRO-RIDE			
PAR PHARM	EQ 5MG ANHYDROUS; 50MG	A070347 001	Dec 25, 1990
MODURETIC 5-50			
+	MERCK	EQ 5MG ANHYDROUS; 50MG **	N018201 001

AMINO ACIDS

INJECTABLE; INJECTION

AMINESS 5.2% ESSENTIAL AMINO ACIDS W/ HISTADINE

HOSPIRA	5.2% (5.2GM/100ML)	N018901 001	Apr 06, 1984
AMINOSYN 10%			
ICU MEDICAL INC	10% (10GM/100ML)	N017673 003	
AMINOSYN 10% (PH6)			
ICU MEDICAL INC	10% (10GM/100ML)	N017673 008	Nov 18, 1985
AMINOSYN 3.5%			
ICU MEDICAL INC	3.5% (3.5GM/100ML)	N017789 004	
AMINOSYN 3.5% IN PLASTIC CONTAINER			
ABBOTT	3.5% (3.5GM/100ML)	N018804 001	May 15, 1984
	3.5% (3.5GM/100ML)	N018875 001	Aug 08, 1984
AMINOSYN 5%			
ICU MEDICAL INC	5% (5GM/100ML)	N017673 001	
AMINOSYN 7%			
ICU MEDICAL INC	7% (7GM/100ML)	N017673 002	
AMINOSYN 7% (PH6)			
ICU MEDICAL INC	7% (7GM/100ML)	N017673 006	Nov 18, 1985
AMINOSYN 8.5%			
ICU MEDICAL INC	8.5% (8.5GM/100ML)	N017673 004	
AMINOSYN 8.5% (PH6)			
ICU MEDICAL INC	8.5% (8.5GM/100ML)	N017673 007	Nov 18, 1985
AMINOSYN II 10%			
ICU MEDICAL INC	10% (10GM/100ML)	N019438 005	Apr 03, 1986
AMINOSYN II 3.5%			
ICU MEDICAL INC	3.5% (3.5GM/100ML)	N019438 001	Apr 03, 1986
AMINOSYN II 3.5% IN PLASTIC CONTAINER			
ABBOTT	3.5% (3.5GM/100ML)	N019491 001	Oct 10, 1986
AMINOSYN II 5%			
ICU MEDICAL INC	5% (5GM/100ML)	N019438 002	Apr 03, 1986
AMINOSYN II 7%			
ICU MEDICAL INC	7% (7GM/100ML)	N019438 003	Apr 03, 1986
AMINOSYN II 8.5%			
ICU MEDICAL INC	8.5% (8.5GM/100ML)	N019438 004	Apr 03, 1986
AMINOSYN-HBC 7%			
ICU MEDICAL INC	7% (7GM/100ML)	N019374 001	Jul 12, 1985
AMINOSYN-HBC 7% IN PLASTIC CONTAINER			
ABBOTT	7% (7GM/100ML)	N019400 001	Jul 23, 1986
AMINOSYN-HF 8%			
ICU MEDICAL INC	8% (8GM/100ML)	A020345 001	Apr 04, 1996

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

AMINO ACIDS

INJECTABLE; INJECTION

AMINOSYN-RF 5.2%				
ICU MEDICAL INC	5.2% (5.2GM/100ML)		N018429 001	
BRANCHAMIN 4%				
BAXTER HLTHCARE	4% (4GM/100ML)		N018678 001	Sep 28, 1984
BRANCHAMIN 4% IN PLASTIC CONTAINER				
BAXTER HLTHCARE	4% (4GM/100ML)		N018684 001	Sep 28, 1984
FREAMINE 8.5%				
B BRAUN	8.5% (8.5GM/100ML)		N016822 001	
FREAMINE HBC 6.9%				
B BRAUN	6.9% (6.9GM/100ML)		N016822 006	May 17, 1983
FREAMINE II 8.5%				
B BRAUN	8.5% (8.5GM/100ML)		N016822 002	
FREAMINE III 10%				
B BRAUN	10% (10GM/100ML)		N016822 005	
FREAMINE III 8.5%				
B BRAUN	8.5% (8.5GM/100ML)		N016822 004	
HEPATAMINE 8%				
B BRAUN	8% (8GM/100ML)		N018676 001	Aug 03, 1982
HEPATASOL 8%				
BAXTER HLTHCARE	8% (8GM/100ML)		A020360 001	Apr 04, 1996
NEOPHAM 6.4%				
HOSPIRA	6.4% (6.4GM/100ML)		N018792 001	Jan 17, 1984
NEPHRAMINE 5.4%				
B BRAUN	5.4% (5.4GM/100ML)		N017766 001	
NOVAMINE 11.4%				
HOSPIRA INC	11.4% (11.4GM/100ML)		N017957 003	Aug 09, 1982
NOVAMINE 15%				
HOSPIRA INC	15% (75GM/500ML)		N017957 004	Nov 28, 1986
NOVAMINE 15% SULFITE FREE IN PLASTIC CONTAINER				
BAXTER HLTHCARE	15% (15GM/100ML) **		N020107 001	Feb 05, 1993
NOVAMINE 8.5%				
HOSPIRA INC	8.5% (8.5GM/100ML)		N017957 002	Aug 09, 1982
RENAMIN W/O ELECTROLYTES				
BAXTER HLTHCARE	6.5% (6.5GM/100ML)		N017493 007	Oct 15, 1982
TRAVASOL 10% W/O ELECTROLYTES				
BAXTER HLTHCARE	10% (10GM/100ML)		N017493 006	
TRAVASOL 5.5% W/O ELECTROLYTES				
BAXTER HLTHCARE	5.5% (5.5GM/100ML)		N017493 004	
TRAVASOL 8.5% W/O ELECTROLYTES				
BAXTER HLTHCARE	8.5% (8.5GM/100ML)		N017493 005	

AMINO ACIDS; CALCIUM ACETATE; GLYCERIN; MAGNESIUM ACETATE; PHOSPHORIC ACID; POTASSIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE

INJECTABLE; INJECTION

PROCALAMINE

B BRAUN	3%;26MG/100ML;3GM/100ML;54MG/100ML;41MG/100ML;150MG/100ML;200MG/100ML;120MG/100ML		N018582 001	May 08, 1982
---------	---	--	-------------	--------------

AMINO ACIDS; CALCIUM CHLORIDE; DEXTROSE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; POTASSIUM PHOSPHATE, DIBASIC; SODIUM CHLORIDE

INJECTABLE; INJECTION

AMINOSYN II 3.5% W/ ELECTROLYTES IN DEXTROSE 25% W/ CALCIUM IN PLASTIC CONTAINER				
ABBOTT	3.5%;36.8MG/100ML;25GM/100ML;51MG/100ML;22.4MG/100ML;261MG/100ML;205MG/100ML		N019714 001	Sep 12, 1988
HOSPIRA INC	3.5%;36.8MG/100ML;25GM/100ML;51MG/100ML;22.4MG/100ML;261MG/100ML;205MG/100ML		N019683 001	Nov 07, 1988
AMINOSYN II 4.25% W/ ELECTROLYTES IN DEXTROSE 20% W/ CALCIUM IN PLASTIC CONTAINER				
ABBOTT	4.25%;36.8MG/100ML;20GM/100ML;51MG/100ML;22.4MG/100ML;261MG/100ML;205MG/100ML		N019714 002	Sep 12, 1988
HOSPIRA INC	4.25%;36.8MG/100ML;20GM/100ML;51MG/100ML;22.4MG/100ML;261MG/100ML;205MG/100ML		N019683 002	Nov 07, 1988
AMINOSYN II 4.25% W/ ELECTROLYTES IN DEXTROSE 25% W/ CALCIUM IN PLASTIC CONTAINER				
ABBOTT	4.25%;36.8MG/100ML;25GM/100ML;51MG/100ML;22.4MG/100ML;261MG/100ML;205MG/100ML		N019714 004	Sep 12, 1988

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

AMINO ACIDS; CALCIUM CHLORIDE; DEXTROSE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; POTASSIUM PHOSPHATE, DIBASIC; SODIUM CHLORIDE

INJECTABLE; INJECTION

AMINOSYN II 4.25% W/ ELECTROLYTES IN DEXTROSE 25% W/ CALCIUM IN PLASTIC CONTAINER			
HOSPIRA INC	4.25%;36.8MG/100ML;25GM/100ML;51MG/100ML L;22.4MG/100ML;261MG/100ML;205MG/100ML	N019683 003	Nov 07, 1988
AMINOSYN II 5% W/ ELECTROLYTES IN DEXTROSE 25% W/ CALCIUM IN PLASTIC CONTAINER			
ABBOTT	5%;36.8MG/100ML;25GM/100ML;51MG/100ML;2 2.4MG/100ML;261MG/100ML;205MG/100ML	N019714 003	Sep 12, 1988
HOSPIRA INC	5%;36.8MG/100ML;25GM/100ML;51MG/100ML;2 2.4MG/100ML;261MG/100ML;205MG/100ML	N019683 004	Nov 07, 1988

AMINO ACIDS; DEXTROSE

INJECTABLE; INJECTION

AMINOSYN 3.5% W/ DEXTROSE 25% IN PLASTIC CONTAINER			
ABBOTT	3.5%;25GM/100ML	N019118 001	Oct 11, 1984
AMINOSYN 3.5% W/ DEXTROSE 5% IN PLASTIC CONTAINER			
ABBOTT	3.5%;5GM/100ML	N019120 001	Oct 11, 1984
AMINOSYN 4.25% W/ DEXTROSE 25% IN PLASTIC CONTAINER			
ABBOTT	4.25%;25GM/100ML	N019119 001	Oct 11, 1984
AMINOSYN II 3.5% IN DEXTROSE 25% IN PLASTIC CONTAINER			
ABBOTT	3.5%;25GM/100ML	N019505 002	Nov 07, 1986
	3.5%;25GM/100ML	N019713 006	Sep 09, 1988
HOSPIRA	3.5%;25GM/100ML	N019681 001	Nov 01, 1988
AMINOSYN II 3.5% IN DEXTROSE 5% IN PLASTIC CONTAINER			
ABBOTT	3.5%;5GM/100ML	N019506 001	Nov 07, 1986
	3.5%;5GM/100ML	N019713 002	Sep 09, 1988
HOSPIRA	3.5%;5GM/100ML	N019681 002	Nov 01, 1988
AMINOSYN II 4.25% IN DEXTROSE 10% IN PLASTIC CONTAINER			
ABBOTT	4.25%;10GM/100ML	N019713 001	Sep 09, 1988
HOSPIRA	4.25%;10GM/100ML	N019681 004	Nov 01, 1988
AMINOSYN II 4.25% IN DEXTROSE 20% IN PLASTIC CONTAINER			
ABBOTT	4.25%;20GM/100ML	N019713 004	Sep 09, 1988
HOSPIRA	4.25%;20GM/100ML	N019681 005	Nov 01, 1988
AMINOSYN II 4.25% IN DEXTROSE 25% IN PLASTIC CONTAINER			
ABBOTT	4.25%;25GM/100ML	N019504 002	Nov 07, 1986
	4.25%;25GM/100ML	N019713 005	Sep 09, 1988
HOSPIRA	4.25%;25GM/100ML	N019681 003	Nov 01, 1988
AMINOSYN II 5% IN DEXTROSE 25% IN PLASTIC CONTAINER			
ABBOTT	5%;25GM/100ML	N019565 001	Dec 17, 1986
	5%;25GM/100ML	N019713 003	Sep 09, 1988
HOSPIRA	5%;25GM/100ML	N019681 006	Nov 01, 1988
TRAVASOL 2.75% IN DEXTROSE 10% IN PLASTIC CONTAINER			
BAXTER HLTHCARE	2.75%;10GM/100ML	N019520 002	Sep 23, 1988
TRAVASOL 2.75% IN DEXTROSE 15% IN PLASTIC CONTAINER			
BAXTER HLTHCARE	2.75%;15GM/100ML	N019520 003	Sep 23, 1988
TRAVASOL 2.75% IN DEXTROSE 20% IN PLASTIC CONTAINER			
BAXTER HLTHCARE	2.75%;20GM/100ML	N019520 004	Sep 23, 1988
TRAVASOL 2.75% IN DEXTROSE 25% IN PLASTIC CONTAINER			
BAXTER HLTHCARE	2.75%;25GM/100ML	N019520 005	Sep 23, 1988
TRAVASOL 2.75% IN DEXTROSE 5% IN PLASTIC CONTAINER			
BAXTER HLTHCARE	2.75%;5GM/100ML	N019520 001	Sep 23, 1988
TRAVASOL 4.25% IN DEXTROSE 10% IN PLASTIC CONTAINER			
BAXTER HLTHCARE	4.25%;10GM/100ML	N019520 007	Sep 23, 1988
TRAVASOL 4.25% IN DEXTROSE 15% IN PLASTIC CONTAINER			
BAXTER HLTHCARE	4.25%;15GM/100ML	N019520 008	Sep 23, 1988
TRAVASOL 4.25% IN DEXTROSE 20% IN PLASTIC CONTAINER			
BAXTER HLTHCARE	4.25%;20GM/100ML	N019520 009	Sep 23, 1988
TRAVASOL 4.25% IN DEXTROSE 25% IN PLASTIC CONTAINER			
BAXTER HLTHCARE	4.25%;25GM/100ML	N019520 010	Sep 23, 1988
TRAVASOL 4.25% IN DEXTROSE 5% IN PLASTIC CONTAINER			
BAXTER HLTHCARE	4.25%;5GM/100ML	N019520 006	Sep 23, 1988

AMINO ACIDS; DEXTROSE; MAGNESIUM CHLORIDE; POTASSIUM ACETATE; POTASSIUM CHLORIDE; POTASSIUM PHOSPHATE, DIBASIC; SODIUM CHLORIDE

INJECTABLE; INJECTION

AMINOSYN II 4.25% W/ ELECT AND ADJUSTED PHOSPHATE IN DEXTROSE 10% IN PLASTIC CONTAINER			
ABBOTT	4.25%;10GM/100ML;51MG/100ML;176.5MG/100 ML;22.4MG/100ML;104.5MG/100ML;205MG/100 ML	N019712 002	Sep 08, 1988
HOSPIRA INC	4.25%;10GM/100ML;51MG/100ML;176.5	N019682 003	Nov 01, 1988

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

AMINO ACIDS; DEXTROSE; MAGNESIUM CHLORIDE; POTASSIUM ACETATE; POTASSIUM CHLORIDE; POTASSIUM PHOSPHATE, DIBASIC; SODIUM CHLORIDE

INJECTABLE; INJECTION

AMINOSYN II 4.25% W/ ELECT AND ADJUSTED PHOSPHATE IN DEXTROSE 10% IN PLASTIC CONTAINER
 MG/100ML; 22.4MG/100ML; 104.5MG/100ML; 205
 MG/100ML

AMINO ACIDS; DEXTROSE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; POTASSIUM PHOSPHATE, DIBASIC; SODIUM CHLORIDE

INJECTABLE; INJECTION

AMINOSYN II 3.5% W/ ELECTROLYTES IN DEXTROSE 25% IN PLASTIC CONTAINER
 ABBOTT 3.5%; 25GM/100ML; 51MG/100ML; 22.4MG/100ML N019564 002 Dec 16, 1986
 ; 261MG/100ML; 205MG/100ML

AMINOSYN II 4.25% W/ ELECTROLYTES IN DEXTROSE 25% IN PLASTIC CONTAINER
 ABBOTT 4.25%; 25GM/100ML; 51MG/100ML; 22.4MG/100ML N019564 004 Dec 16, 1986
 ; 261MG/100ML; 205MG/100ML

AMINO ACIDS; DEXTROSE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM CHLORIDE; SODIUM PHOSPHATE, DIBASIC, HEPTAHYDRATE

INJECTABLE; INJECTION

AMINOSYN II 3.5% M IN DEXTROSE 5% IN PLASTIC CONTAINER
 ABBOTT 3.5%; 5GM/100ML; 30MG/100ML; 97MG/100ML; 12 N019564 001 Dec 16, 1986
 0MG/100ML; 49.3MG/100ML

3.5%; 5GM/100ML; 30MG/100ML; 97MG/100ML; 12 N019712 001 Sep 08, 1988
 0MG/100ML; 49.3MG/100ML

HOSPIRA INC 3.5%; 5GM/100ML; 30MG/100ML; 97MG/100ML; 12 N019682 001 Nov 01, 1988
 0MG/100ML; 49.3MG/100ML

AMINOSYN II 4.25% M IN DEXTROSE 10% IN PLASTIC CONTAINER
 ABBOTT 4.25%; 10GM/100ML; 30MG/100ML; 97MG/100ML; N019564 003 Dec 16, 1986
 120MG/100ML; 49.3MG/100ML

HOSPIRA INC 4.25%; 5GM/100ML; 30MG/100ML; 97MG/100ML; 1 N019682 002 Nov 01, 1988
 20MG/100ML; 49.3MG/100ML

AMINO ACIDS; DEXTROSE; MAGNESIUM CHLORIDE; POTASSIUM PHOSPHATE, DIBASIC; SODIUM ACETATE; SODIUM CHLORIDE

INJECTABLE; INJECTION

TRAVASOL 2.75% SULFITE FREE W/ ELECTROLYTES IN DEXTROSE 10% IN PLASTIC CONTAINER
 BAXTER HLTHCARE 2.75%; 10GM/100ML; 51MG/100ML; 261MG/100ML N020147 002 Oct 23, 1995
 ; 216MG/100ML; 112MG/100ML

TRAVASOL 2.75% SULFITE FREE W/ ELECTROLYTES IN DEXTROSE 15% IN PLASTIC CONTAINER
 BAXTER HLTHCARE 2.75%; 15GM/100ML; 51MG/100ML; 261MG/100ML N020147 003 Oct 23, 1995
 ; 216MG/100ML; 112MG/100ML

TRAVASOL 2.75% SULFITE FREE W/ ELECTROLYTES IN DEXTROSE 20% IN PLASTIC CONTAINER
 BAXTER HLTHCARE 2.75%; 20GM/100ML; 51MG/100ML; 261MG/100ML N020147 004 Oct 23, 1995
 ; 216MG/100ML; 112MG/100ML

TRAVASOL 2.75% SULFITE FREE W/ ELECTROLYTES IN DEXTROSE 25% IN PLASTIC CONTAINER
 BAXTER HLTHCARE 2.75%; 25GM/100ML; 51MG/100ML; 261MG/100ML N020147 005 Oct 23, 1995
 ; 216MG/100ML; 112MG/100ML

TRAVASOL 2.75% SULFITE FREE W/ ELECTROLYTES IN DEXTROSE 5% IN PLASTIC CONTAINER
 BAXTER HLTHCARE 2.75%; 5GM/100ML; 51MG/100ML; 261MG/100ML; N020147 001 Oct 23, 1995
 216MG/100ML; 112MG/100ML

TRAVASOL 4.25% SULFITE FREE W/ ELECTROLYTES IN DEXTROSE 10% IN PLASTIC CONTAINER
 BAXTER HLTHCARE 4.25%; 10GM/100ML; 51MG/100ML; 261MG/100ML N020147 007 Oct 23, 1995
 ; 297MG/100ML; 77MG/100ML

TRAVASOL 4.25% SULFITE FREE W/ ELECTROLYTES IN DEXTROSE 15% IN PLASTIC CONTAINER
 BAXTER HLTHCARE 4.25%; 15GM/100ML; 51MG/100ML; 261MG/100ML N020147 008 Oct 23, 1995
 ; 297MG/100ML; 77MG/100ML

TRAVASOL 4.25% SULFITE FREE W/ ELECTROLYTES IN DEXTROSE 20% IN PLASTIC CONTAINER
 BAXTER HLTHCARE 4.25%; 20GM/100ML; 51MG/100ML; 261MG/100ML N020147 009 Oct 23, 1995
 ; 297MG/100ML; 77MG/100ML

TRAVASOL 4.25% SULFITE FREE W/ ELECTROLYTES IN DEXTROSE 25% IN PLASTIC CONTAINER
 BAXTER HLTHCARE 4.25%; 25GM/100ML; 51MG/100ML; 261MG/100ML N020147 010 Oct 23, 1995
 ; 297MG/100ML; 77MG/100ML

TRAVASOL 4.25% SULFITE FREE W/ ELECTROLYTES IN DEXTROSE 5% IN PLASTIC CONTAINER
 BAXTER HLTHCARE 4.25%; 5GM/100ML; 51MG/100ML; 261MG/100ML; N020147 006 Oct 23, 1995
 297MG/100ML; 77MG/100ML

AMINO ACIDS; MAGNESIUM ACETATE; PHOSPHORIC ACID; POTASSIUM ACETATE; POTASSIUM CHLORIDE; SODIUM ACETATE

INJECTABLE; INJECTION

FREAMINE III 8.5% W/ ELECTROLYTES
 B BRAUN 8.5%; 110MG/100ML; 230MG/100ML; 10MG/100ML N016822 007 Jul 01, 1988
 ; 440MG/100ML; 690MG/100ML

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

AMINO ACIDS; MAGNESIUM ACETATE; PHOSPHORIC ACID; POTASSIUM ACETATE; SODIUM CHLORIDE

INJECTABLE; INJECTION

AMINOSYN 3.5% M

ICU MEDICAL INC	3.5%; 21MG/100ML; 40MG/100ML; 128MG/100ML; 234MG/100ML	N017789 003
-----------------	---	-------------

AMINOSYN 3.5% M IN PLASTIC CONTAINER

ABBOTT	3.5%; 21MG/100ML; 40MG/100ML; 128MG/100ML; 234MG/100ML	N018804 002	May 15, 1984
	3.5%; 21MG/100ML; 40MG/100ML; 128MG/100ML; 234MG/100ML	N018875 002	Aug 08, 1984

AMINO ACIDS; MAGNESIUM ACETATE; PHOSPHORIC ACID; POTASSIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE

INJECTABLE; INJECTION

FREAMINE III 3% W/ ELECTROLYTES

B BRAUN	3%; 54MG/100ML; 40MG/100ML; 150MG/100ML; 20 0MG/100ML; 120MG/100ML	N016822 003
---------	---	-------------

AMINO ACIDS; MAGNESIUM ACETATE; POTASSIUM ACETATE; SODIUM CHLORIDE

INJECTABLE; INJECTION

AMINOSYN 3.5% M

ICU MEDICAL INC	3.5%; 21MG/100ML; 128MG/100ML; 234MG/100ML	N017789 005
-----------------	--	-------------

AMINO ACIDS; MAGNESIUM ACETATE; POTASSIUM ACETATE; SODIUM CHLORIDE; SODIUM PHOSPHATE, DIBASIC, HEPTAHYDRATE

INJECTABLE; INJECTION

AMINOSYN II 3.5% M IN PLASTIC CONTAINER

ABBOTT	3.5%; 32MG/100ML; 128MG/100ML; 222MG/100ML ; 49MG/100ML	N019493 001	Oct 16, 1986
--------	--	-------------	--------------

AMINO ACIDS; MAGNESIUM CHLORIDE; POTASSIUM ACETATE; POTASSIUM CHLORIDE; SODIUM ACETATE

INJECTABLE; INJECTION

VEINAMINE 8%

HOSPIRA INC	8%; 61MG/100ML; 211MG/100ML; 56MG/100ML; 38 8MG/100ML	N017957 001
-------------	--	-------------

AMINO ACIDS; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; POTASSIUM PHOSPHATE, DIBASIC; SODIUM CHLORIDE

INJECTABLE; INJECTION

AMINOSYN II 10% W/ ELECTROLYTES

ICU MEDICAL INC	10%; 102MG/100ML; 45MG/100ML; 522MG/100ML; 410MG/100ML	N019437 004	Apr 03, 1986
-----------------	---	-------------	--------------

AMINOSYN II 7% W/ ELECTROLYTES

ICU MEDICAL INC	7%; 102MG/100ML; 45MG/100ML; 522MG/100ML; 4 10MG/100ML	N019437 006	Apr 03, 1986
-----------------	---	-------------	--------------

AMINOSYN II 8.5% W/ ELECTROLYTES

ICU MEDICAL INC	8.5%; 102MG/100ML; 45MG/100ML; 522MG/100ML ; 410MG/100ML	N019437 005	Apr 03, 1986
-----------------	---	-------------	--------------

AMINO ACIDS; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM CHLORIDE; SODIUM PHOSPHATE, DIBASIC

INJECTABLE; INJECTION

AMINOSYN 8.5% W/ELECTROLYTES

ICU MEDICAL INC	8.5%; 102MG/100ML; 487MG/100ML; 28MG/100ML ; 425MG/100ML	N017673 009	Oct 25, 2002
-----------------	---	-------------	--------------

AMINOSYN II 8.5% W/ELECTROLYTES

ICU MEDICAL INC	8.5%; 102MG/100ML; 492MG/100ML; 60MG/100ML ; 425MG/100ML	N019437 008	Oct 25, 2002
-----------------	---	-------------	--------------

AMINO ACIDS; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM CHLORIDE; SODIUM PHOSPHATE, DIBASIC, HEPTAHYDRATE

INJECTABLE; INJECTION

AMINOSYN II 3.5% M

ICU MEDICAL INC	3.5%; 30MG/100ML; 97MG/100ML; 120MG/100ML; 49MG/100ML	N019437 007	Apr 03, 1986
-----------------	--	-------------	--------------

AMINO ACIDS; MAGNESIUM CHLORIDE; POTASSIUM PHOSPHATE, DIBASIC; SODIUM ACETATE; SODIUM CHLORIDE

INJECTABLE; INJECTION

TRAVASOL 3.5% SULFITE FREE W/ ELECTROLYTES IN PLASTIC CONTAINER

BAXTER HLTHCARE	3.5%; 51MG/100ML; 131MG/100ML; 218MG/100ML ; 35MG/100ML	N020177 001	Oct 23, 1995
-----------------	--	-------------	--------------

TRAVASOL 3.5% W/ ELECTROLYTES

BAXTER HLTHCARE	3.5%; 51MG/100ML; 131MG/100ML; 218MG/100ML ; 35MG/100ML	N017493 003
-----------------	--	-------------

TRAVASOL 5.5% SULFITE FREE W/ ELECTROLYTES IN PLASTIC CONTAINER

BAXTER HLTHCARE	5.5%; 102MG/100ML; 522MG/100ML; 431MG/100M L; 224MG/100ML	N020173 001	Oct 27, 1995
-----------------	--	-------------	--------------

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

AMINO ACIDS; MAGNESIUM CHLORIDE; POTASSIUM PHOSPHATE, DIBASIC; SODIUM ACETATE; SODIUM CHLORIDE

INJECTABLE; INJECTION

TRAVASOL 5.5% W/ ELECTROLYTES

BAXTER HLTHCARE	5.5%;102MG/100ML;522MG/100ML;431MG/100ML	N017493 001	
	L;224MG/100ML		

TRAVASOL 8.5% SULFITE FREE W/ ELECTROLYTES IN PLASTIC CONTAINER

BAXTER HLTHCARE	8.5%;102MG/100ML;522MG/100ML;594MG/100ML	N020173 002	Oct 27, 1995
	L;154MG/100ML		

TRAVASOL 8.5% W/ ELECTROLYTES

BAXTER HLTHCARE	8.5%;102MG/100ML;522MG/100ML;594MG/100ML	N017493 002	
	L;154MG/100ML		

AMINO ACIDS; MAGNESIUM CHLORIDE; POTASSIUM PHOSPHATE, DIBASIC; SODIUM CHLORIDE

INJECTABLE; INJECTION

AMINOSYN 7% W/ ELECTROLYTES

ICU MEDICAL INC	7%;102MG/100ML;522MG/100ML;410MG/100ML	N017789 002	
-----------------	--	-------------	--

AMINOSYN 8.5% W/ ELECTROLYTES

ICU MEDICAL INC	8.5%;102MG/100ML;522MG/100ML;410MG/100ML	N017673 005	
	L		

AMINOCAPROIC ACID

INJECTABLE; INJECTION

AMICAR

+ EPIC PHARMA LLC	250MG/ML **	N015229 002	
-------------------	-------------	-------------	--

AMINOCAPROIC ACID

ABRAXIS PHARM	250MG/ML	A070522 001	Jun 17, 1986
BAXTER HLTHCARE	250MG/ML	N018590 001	Oct 29, 1982
HOSPIRA	250MG/ML	A070888 001	Jun 16, 1988

SOLUTION; ORAL

AMINOCAPROIC ACID

EPIC PHARMA LLC	0.25GM/ML	A074759 001	Sep 02, 1998
VISTAPHARM LLC	0.25GM/ML	A212814 001	Feb 26, 2020

TABLET; ORAL

AMINOCAPROIC

HIKMA	500MG	A075602 001	May 24, 2001
-------	-------	-------------	--------------

AMINOCAPROIC ACID

ADAPTIS	500MG	A212110 001	Jun 14, 2021
ANI PHARMS	500MG	A211629 001	Dec 14, 2020

AMINOGLUTETHIMIDE

TABLET; ORAL

CYTADREN

NOVARTIS	250MG	N018202 001	
----------	-------	-------------	--

AMINOHIPPURATE SODIUM

INJECTABLE; INJECTION

AMINOHIPPURATE SODIUM

MERCK	20%	N005619 001	
-------	-----	-------------	--

AMINOPHYLLINE

ENEMA; RECTAL

SOMOPHYLLIN

FISONS	300MG/5ML	N018232 001	Apr 02, 1982
--------	-----------	-------------	--------------

INJECTABLE; INJECTION

AMINOPHYLLIN

+ GD SEARLE LLC	25MG/ML	A087243 001	May 24, 1982
-----------------	---------	-------------	--------------

+	25MG/ML	A087621 001	May 24, 1982
---	---------	-------------	--------------

AMINOPHYLLINE

ABRAXIS PHARM	25MG/ML	A084568 001	
	25MG/ML	A087200 001	
	25MG/ML	A087250 001	Jan 06, 1982
	25MG/ML	A087886 001	Aug 30, 1983
	25MG/ML	A088407 001	Jan 25, 1984
AM REGENT	25MG/ML	A087600 001	
ELKINS SINN	25MG/ML	A087239 001	
HOSPIRA	25MG/ML	A087601 001	Jul 23, 1982
INTL MEDICATION	25MG/ML	A087209 001	Feb 01, 1982
	25MG/ML	A087867 001	Nov 10, 1983
	25MG/ML	A087868 001	Nov 10, 1983
KING PHARMS	25MG/ML	A086606 001	
LUITPOLD	25MG/ML	A087240 001	
LYPHOMED	25MG/ML	A087431 001	

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

AMINOPHYLLINE

INJECTABLE; INJECTION

AMINOPHYLLINE

PHARMA SERVE NY	25MG/ML	A087387 001	Jun 03, 1983
	25MG/ML	A087392 001	Dec 15, 1983
SMITH AND NEPHEW	25MG/ML	A088429 001	May 30, 1985
	25MG/ML	A088749 001	May 30, 1985
TEVA PARENTERAL	25MG/ML	A081142 001	Sep 25, 1991
AMINOPHYLLINE IN SODIUM CHLORIDE 0.45%			
HOSPIRA	100MG/100ML	A088147 002	May 03, 1983
	200MG/100ML	A088147 003	May 03, 1983
AMINOPHYLLINE IN SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER			
HOSPIRA	100MG/100ML	N018924 001	Dec 12, 1984
	200MG/100ML	N018924 002	Dec 12, 1984
	400MG/100ML	N018924 003	Dec 12, 1984
	500MG/100ML	N018924 004	Dec 12, 1984

SOLUTION; ORAL

AMINOPHYLLINE

MORTON GROVE	105MG/5ML	A088156 001	Dec 05, 1983
ROXANE	105MG/5ML	A088126 001	Aug 19, 1983
AMINOPHYLLINE DYE FREE			
ACTAVIS MID ATLANTIC	105MG/5ML	A087727 001	Apr 16, 1982
SOMOPHYLLIN			
FISONS	105MG/5ML	A086466 001	
SOMOPHYLLIN-DF			
FISONS	105MG/5ML	A087045 001	
SUPPOSITORY; RECTAL			
TRUPHYLLINE			
COSETTE	250MG	A085498 001	Mar 23, 1983
	500MG	A085498 002	Jan 03, 1983

TABLET; ORAL

AMINOPHYLLIN

GD SEARLE LLC	100MG	N002386 002	
	200MG	N002386 003	
AMINOPHYLLINE			
ANI PHARMS	100MG	A085261 004	
	200MG	A085261 002	
ASCOT	100MG	A087522 001	Feb 12, 1982
	200MG	A087523 001	Feb 12, 1982
BARR	100MG	A088297 001	Aug 19, 1983
	200MG	A088298 001	Aug 19, 1983
CHARTWELL MOLECULAR	100MG	A084588 001	
	200MG	A084588 002	
DURAMED PHARMS BARR	100MG	A088182 001	Mar 31, 1983
	200MG	A088183 001	Mar 31, 1983
HALSEY	100MG	A084674 001	
HIKMA INTL PHARMS	100MG	A084540 001	
	200MG	A085003 001	
IMPAX LABS	100MG	A084574 001	
	200MG	A084576 001	
KV PHARM	100MG	A085284 001	
	200MG	A085289 001	
PAL PAK	100MG	A084533 001	
PANRAY	100MG	A084552 001	
	200MG	A084552 002	
PUREPAC PHARM	100MG	A084699 001	
	200MG	A085333 001	
ROXANE	100MG	A087500 001	Feb 09, 1982
	200MG	A087501 001	Feb 09, 1982
VALEANT PHARM INTL	200MG	A084563 001	
VANGARD	100MG	A088314 001	Oct 03, 1983
	200MG	A088319 001	Oct 03, 1983
VINTAGE PHARMS	100MG	A085409 001	
	200MG	A085410 001	
WATSON LABS	100MG	A085567 001	
	200MG	A085564 001	

TABLET, DELAYED RELEASE; ORAL

AMINOPHYLLINE

IMPAX LABS	100MG	A084577 001	
------------	-------	-------------	--

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

AMINOPHYLLINE

TABLET, DELAYED RELEASE;ORAL
AMINOPHYLLINE

	200MG	A084575 001
TABLICAPS	100MG	A084632 002
VALE	100MG	A084531 001
	200MG	A084530 001

TABLET, EXTENDED RELEASE;ORAL
PHYLLOCONTIN

PHARM RES ASSOC	225MG	A086760 001
-----------------	-------	-------------

AMINOSALICYLATE SODIUM

POWDER;ORAL

P.A.S. SODIUM

CENTURY PHARMS	4GM/PACKET	A080947 001
----------------	------------	-------------

SODIUM AMINOSALICYLATE

HEXCEL	100%	A080097 001
--------	------	-------------

TABLET;ORAL

PARASAL SODIUM

PANRAY	500MG	N006811 006
	1GM	N006811 011

SODIUM P.A.S.

LANNETT	500MG	A080138 002
---------	-------	-------------

TEEBACIN

CONSOLIDATED MIDLAND	500MG	N007320 002
----------------------	-------	-------------

AMINOSALICYLATE SODIUM; AMINOSALICYLIC ACID

TABLET;ORAL

NEOPASALATE

MEDPOINTE PHARM HLC	846MG;112MG	A080059 002
---------------------	-------------	-------------

AMINOSALICYLIC ACID

TABLET;ORAL

PARASAL

PANRAY	500MG	N006811 001
	1GM	N006811 002

AMINOSALICYLIC ACID RESIN COMPLEX

POWDER;ORAL

REZIPAS

BRISTOL MYERS SQUIBB	EQ 500MG BASE/GM	N009052 001
----------------------	------------------	-------------

AMIODARONE HYDROCHLORIDE

INJECTABLE; INJECTION

AMIODARONE HYDROCHLORIDE

BEDFORD	50MG/ML	A076018 001	Oct 15, 2002
BEDFORD LABS	50MG/ML	A076299 001	Oct 24, 2002
BEN VENUE	50MG/ML	A076088 001	Oct 15, 2002
DR REDDYS	50MG/ML	A076163 001	Sep 05, 2003
EPIC PHARMA LLC	50MG/ML	A076232 001	Jul 05, 2006
EUGIA PHARMA	50MG/ML	A204550 001	Oct 25, 2017
HOSPIRA	50MG/ML	A075955 001	Oct 18, 2002
	50MG/ML	A076108 001	Oct 15, 2002
	50MG/ML	A203884 001	Nov 25, 2013
	50MG/ML	A203885 001	Nov 25, 2013
INTL MEDICATION SYS	50MG/ML	N021594 001	Feb 04, 2004
PAR STERILE PRODUCTS	50MG/ML	A076394 001	Apr 25, 2003

CORDARONE

+ WYETH PHARMS INC

50MG/ML **	N020377 001	Aug 03, 1995
------------	-------------	--------------

NEXTERONE

+ BAXTER HLTHCARE

50MG/ML **	N022325 001	Dec 24, 2008
------------	-------------	--------------

TABLET;ORAL

AMIODARONE HYDROCHLORIDE

NORVIUM BIOSCIENCE	200MG	A075188 001	Feb 24, 1999
TEVA	200MG	A074895 001	Apr 16, 1999
UPSHER SMITH LABS	200MG	A075315 001	Dec 23, 1998
	400MG	A075315 002	Jun 30, 2000

CORDARONE

+ WYETH PHARMS

200MG **	N018972 001	Dec 24, 1985
----------	-------------	--------------

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

AMITRIPTYLINE HYDROCHLORIDE

CONCENTRATE; ORAL

ENDEP

+ ROCHE 40MG/ML ** A085749 001

INJECTABLE; INJECTION

AMITRIPTYLINE HYDROCHLORIDE

WATSON LABS 10MG/ML A085594 001

ELAVIL

ASTRAZENECA 10MG/ML N012704 001

TABLET; ORAL

AMITID

BRISTOL MYERS SQUIBB 10MG A086454 001

25MG A086454 002

50MG A086454 003

75MG A086454 004

100MG A086454 005

AMITRIL

WARNER CHILCOTT 10MG A083939 001

25MG A083937 001

50MG A083938 002

75MG A084957 001

100MG A085093 001

150MG A086295 001

AMITRIPTYLINE HYDROCHLORIDE

AM THERAP 25MG A088672 001 Nov 20, 1984

50MG A088673 001 Nov 20, 1984

75MG A088674 001 Nov 20, 1984

100MG A088675 001 Nov 20, 1984

ANI PHARMS 10MG A085031 002

25MG A085031 001

50MG A085031 003

75MG A085031 004

AUROBINDO PHARMA USA 10MG A086009 002

25MG A086009 003

50MG A086009 001

75MG A086009 004

100MG A086009 005

150MG A086009 006

COPLEY PHARM 10MG A088421 001 Apr 30, 1984

25MG A088422 001 Apr 30, 1984

50MG A088423 001 Apr 30, 1984

75MG A088424 001 Apr 30, 1984

100MG A088425 001 Apr 30, 1984

150MG A088426 001 Apr 30, 1984

ENDO OPERATIONS 10MG A040218 001 Sep 11, 1997

25MG A040218 002 Sep 11, 1997

50MG A040218 003 Sep 11, 1997

75MG A040218 004 Sep 11, 1997

100MG A040218 005 Sep 11, 1997

150MG A040218 006 Sep 11, 1997

HALSEY 10MG A085923 001

25MG A085922 001

50MG A085925 001

50MG A087557 001 Mar 05, 1982

75MG A085926 001 May 20, 1983

100MG A085927 001 May 20, 1983

LEDERLE 10MG A086744 001

10MG A087366 001 Jan 04, 1982

25MG A086746 001

25MG A087367 001 May 03, 1982

50MG A086743 001

50MG A087181 001 Jan 04, 1982

75MG A086745 001

75MG A087369 001 Jan 04, 1982

100MG A086747 001

100MG A087368 001 May 03, 1982

150MG A087370 001 Jan 04, 1982

MUTUAL PHARM 10MG A085744 001

25MG A085627 001

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

AMITRIPTYLINE HYDROCHLORIDE

TABLET;ORAL

AMITRIPTYLINE HYDROCHLORIDE

	50MG	A085745	001	
	75MG	A085743	001	
	100MG	A085742	002	May 11, 1982
	150MG	A089423	001	Feb 17, 1987
PAR PHARM	10MG	A088697	001	Sep 25, 1984
	25MG	A088698	001	Sep 25, 1984
	50MG	A088699	001	Sep 25, 1984
	75MG	A088700	001	Sep 25, 1984
	100MG	A088701	001	Sep 25, 1984
	150MG	A088702	001	Sep 25, 1984
PLIVA	10MG	A088883	001	Sep 26, 1984
	25MG	A088884	001	Sep 26, 1984
	50MG	A088885	001	Sep 26, 1984
	75MG	A088886	001	Sep 26, 1984
	100MG	A088887	001	Sep 26, 1984
	150MG	A088888	001	Sep 26, 1984
PUREPAC PHARM	10MG	A088075	001	Sep 16, 1983
	10MG	A088084	001	Jul 18, 1983
	25MG	A088076	001	May 20, 1983
	25MG	A088085	001	Jul 18, 1983
	50MG	A088077	001	Sep 16, 1983
	50MG	A088105	001	Jul 18, 1983
	75MG	A088078	001	Sep 16, 1983
	75MG	A088106	001	Jul 18, 1983
	100MG	A088079	001	Sep 16, 1983
	100MG	A088107	001	Jul 18, 1983
ROXANE	10MG	A086002	001	
	10MG	A086144	001	
	25MG	A085944	001	
	25MG	A086145	001	
	50MG	A085945	001	
	50MG	A086143	001	
	75MG	A086004	001	
	75MG	A086147	001	
	100MG	A086003	001	
	100MG	A086146	001	
	150MG	A086090	001	
	150MG	A086148	001	
SUN PHARM INDS INC	10MG	A040816	002	Jun 27, 2008
	25MG	A040816	001	Jun 27, 2008
	50MG	A040816	003	Jun 27, 2008
	75MG	A040816	004	Jun 27, 2008
	100MG	A040816	005	Jun 27, 2008
	150MG	A040816	006	Jun 27, 2008
SUPERPHARM	10MG	A088853	001	Nov 13, 1984
	25MG	A088854	001	Nov 13, 1984
	50MG	A088855	001	Nov 13, 1984
	75MG	A088856	001	Nov 13, 1984
	100MG	A088857	001	Nov 13, 1984
TEVA	10MG	A086610	001	
	25MG	A086859	001	
	50MG	A086857	001	
	75MG	A086860	001	
	100MG	A085836	001	
	100MG	A086854	001	
	150MG	A086853	001	
UCB INC	10MG	A085864	001	
	25MG	A085935	001	
	50MG	A085936	001	
	75MG	A086337	001	
	100MG	A086336	001	
	150MG	A086335	001	
USL PHARMA	25MG	A087775	001	Feb 10, 1982
VANGARD	10MG	A087632	001	Feb 01, 1982
	50MG	A087616	001	Feb 08, 1982
	75MG	A087617	001	Feb 05, 1982

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

AMITRIPTYLINE HYDROCHLORIDE

TABLET;ORAL

AMITRIPTYLINE HYDROCHLORIDE

		100MG	A087639	001	Feb 08, 1982
	WATSON LABS	10MG	A085816	001	
		10MG	A088620	001	Mar 02, 1984
		25MG	A085817	001	
		25MG	A088621	001	Mar 02, 1984
		50MG	A085815	001	
		50MG	A088622	001	Mar 02, 1984
		75MG	A085819	001	
		75MG	A088633	001	Mar 02, 1984
		100MG	A085820	001	
		100MG	A088634	001	Mar 02, 1984
		150MG	A085821	001	
		150MG	A088635	001	Mar 02, 1984
	WEST WARD	10MG	A087647	001	Mar 05, 1982
		25MG	A087278	001	
	ELAVIL				
	+ ASTRAZENECA	10MG **	N012703	001	
		25MG **	N012703	003	
		50MG **	N012703	004	
		75MG **	N012703	005	
		100MG **	N012703	006	
		150MG **	N012703	007	
	ENDEP				
	ROCHE	10MG	A083639	001	
		25MG	A083639	002	
		50MG	A083639	003	
		75MG	A083639	004	
		100MG	A083639	005	
		150MG	A085303	001	

AMITRIPTYLINE HYDROCHLORIDE; CHLORDIAZEPOXIDE

TABLET;ORAL

CHLORDIAZEPOXIDE AND AMITRIPTYLINE HYDROCHLORIDE

	ANDA REPOSITORY	EQ 12.5MG BASE;5MG	A070765	001	Dec 10, 1986
		EQ 25MG BASE;10MG	A070766	001	Dec 10, 1986
	CHARTWELL RX	EQ 12.5MG BASE;5MG	A072277	001	May 09, 1988
		EQ 25MG BASE;10MG	A072278	001	May 09, 1988
	HERITAGE PHARMA	EQ 12.5MG BASE;5MG	A072052	001	Dec 16, 1988
		EQ 25MG BASE;10MG	A072053	001	Dec 16, 1988
	USL PHARMA	EQ 12.5MG BASE;5MG	A070477	001	Jan 12, 1988
		EQ 25MG BASE;10MG	A070478	001	Jan 12, 1988
	LIMBITROL				
	+ CHARTWELL RX	EQ 12.5MG BASE;5MG **	N016949	001	
	LIMBITROL DS				
	+ CHARTWELL RX	EQ 25MG BASE;10MG **	N016949	002	

AMITRIPTYLINE HYDROCHLORIDE; PERPHENAZINE

TABLET;ORAL

ETRAFON 2-10

	SCHERING	10MG;2MG **	N014713	007	
	ETRAFON 2-25				
	SCHERING	25MG;2MG **	N014713	004	
	ETRAFON-A				
	SCHERING	10MG;4MG **	N014713	002	
	ETRAFON-FORTE				
	SCHERING	25MG;4MG **	N014713	006	
	PERPHENAZINE AND AMITRIPTYLINE HYDROCHLORIDE				
	CHARTWELL RX	10MG;2MG	A073007	001	Oct 17, 1991
		10MG;4MG	A073009	001	Oct 17, 1991
		25MG;2MG	A073008	001	Oct 17, 1991
		25MG;4MG	A073010	001	Oct 17, 1991
	FOSUN PHARMA	10MG;2MG	A071062	001	Nov 27, 1987
		10MG;4MG	A071862	001	Dec 21, 1987
		25MG;2MG	A071063	001	Nov 27, 1987
		25MG;4MG	A071064	001	Nov 27, 1987
		50MG;4MG	A071863	001	Dec 21, 1987
	IVAX SUB TEVA PHARMS	10MG;2MG	A070935	001	Sep 11, 1986

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

AMITRIPTYLINE HYDROCHLORIDE; PERPHENAZINE

TABLET; ORAL

PERPHENAZINE AND AMITRIPTYLINE HYDROCHLORIDE

	10MG; 4MG	A070937	001	Sep 11, 1986
	25MG; 2MG	A070936	001	Sep 11, 1986
	25MG; 4MG	A070938	001	Sep 11, 1986
	50MG; 4MG	A070939	001	Sep 12, 1986
PAR PHARM	10MG; 2MG	A070565	001	Sep 11, 1986
	10MG; 4MG	A070620	001	Sep 11, 1986
	25MG; 2MG	A070621	001	Sep 11, 1986
	25MG; 4MG	A070595	001	Sep 11, 1986
	50MG; 4MG	A070574	001	Sep 11, 1986
SUN PHARM INDUSTRIES	10MG; 2MG	A071077	001	Nov 12, 1986
	10MG; 4MG	A071078	001	Nov 12, 1986
	25MG; 2MG	A070297	001	Nov 12, 1986
	25MG; 4MG	A071079	001	Nov 12, 1986
WATSON LABS	10MG; 2MG	A070373	001	Aug 25, 1986
	10MG; 2MG	A072539	001	Feb 15, 1989
	10MG; 4MG	A070375	001	Aug 25, 1986
	10MG; 4MG	A072540	001	Feb 15, 1989
	25MG; 2MG	A070374	001	Aug 25, 1986
	25MG; 2MG	A072541	001	Feb 15, 1989
	25MG; 4MG	A070376	001	Aug 25, 1986
	25MG; 4MG	A072134	001	Feb 15, 1989
	50MG; 4MG	A070377	001	Nov 04, 1986
	50MG; 4MG	A071558	001	Mar 02, 1987
	50MG; 4MG	A072135	001	Feb 15, 1989
TRIAVIL 2-10				
NEW RIVER	10MG; 2MG **	N014715	004	
TRIAVIL 2-25				
NEW RIVER	25MG; 2MG **	N014715	002	
TRIAVIL 4-10				
NEW RIVER	10MG; 4MG **	N014715	003	
TRIAVIL 4-25				
NEW RIVER	25MG; 4MG **	N014715	005	
TRIAVIL 4-50				
NEW RIVER	50MG; 4MG **	N014715	006	

AMLEXANOX

PASTE; DENTAL

APHTHASOL

ULURU

5%

N020511 001 Dec 17, 1996

PATCH; TOPICAL

AMLEXANOX

ULURU

2MG

N021727 001 Sep 29, 2004

AMLODIPINE BENZOATE

SUSPENSION; ORAL

AMLODIPINE BENZOATE

AMNEAL

EQ 1MG BASE/ML

A215035 001 Jun 13, 2023

AMLODIPINE BESYLATE

TABLET; ORAL

AMLODIPINE BESYLATE

AMNEAL PHARMS NY

EQ 2.5MG BASE

A078477 001 Jan 16, 2008

EQ 5MG BASE

A078477 002 Jan 16, 2008

EQ 10MG BASE

A078477 003 Jan 16, 2008

CHARTWELL RX

EQ 2.5MG BASE

A076859 001 Sep 10, 2007

EQ 5MG BASE

A076859 002 Sep 10, 2007

EQ 10MG BASE

A076859 003 Sep 10, 2007

GEDEON RICHTER USA

EQ 2.5MG BASE

A077333 001 Jul 17, 2007

EQ 5MG BASE

A077333 002 Jul 17, 2007

EQ 10MG BASE

A077333 003 Jul 17, 2007

GENPHARM

EQ 2.5MG BASE

A077362 001 Jul 09, 2007

EQ 5MG BASE

A077362 002 Jul 09, 2007

EQ 10MG BASE

A077362 003 Jul 09, 2007

GRAVITI PHARMS

EQ 5MG BASE

A201380 001 Apr 13, 2012

EQ 10MG BASE

A201380 002 Apr 13, 2012

HIKMA

EQ 2.5MG BASE

A077262 001 Jul 09, 2007

EQ 5MG BASE

A077262 002 Jul 09, 2007

EQ 10MG BASE

A077262 003 Jul 09, 2007

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

AMLODIPINE BESYLATE

TABLET;ORAL

AMLODIPINE BESYLATE

HIKMA PHARMS	EQ 2.5MG BASE	A077771 001	Apr 12, 2011
	EQ 5MG BASE	A077771 002	Apr 12, 2011
	EQ 10MG BASE	A077771 003	Apr 12, 2011
MYLAN	EQ 2.5MG BASE	A076418 001	Oct 03, 2005
	EQ 5MG BASE	A076418 002	Oct 03, 2005
	EQ 10MG BASE	A076418 003	Oct 03, 2005
NORVIUM BIOSCIENCE	EQ 2.5MG BASE	A078224 001	Feb 27, 2008
	EQ 5MG BASE	A078224 002	Feb 27, 2008
	EQ 10MG BASE	A078224 003	Feb 27, 2008
PURACAP PHARM	EQ 2.5MG BASE	A078131 001	Sep 04, 2007
	EQ 5MG BASE	A078131 002	Sep 04, 2007
	EQ 10MG BASE	A078131 003	Sep 04, 2007
SCIEGEN PHARMS INC	EQ 2.5MG BASE	A077759 001	Jul 09, 2007
	EQ 5MG BASE	A077759 002	Jul 09, 2007
	EQ 10MG BASE	A077759 003	Jul 09, 2007
SOVEREIGN PHARMS	EQ 2.5MG BASE	A204900 001	Jul 23, 2015
	EQ 5MG BASE	A204900 002	Jul 23, 2015
	EQ 10MG BASE	A204900 003	Jul 23, 2015
SUN PHARM INDS INC	EQ 2.5MG BASE	A078231 001	Nov 30, 2007
	EQ 5MG BASE	A078231 002	Nov 30, 2007
	EQ 10MG BASE	A078231 003	Nov 30, 2007
SUN PHARM INDS LTD	EQ 2.5MG BASE	A077974 001	Jul 09, 2007
	EQ 5MG BASE	A077974 002	Jul 09, 2007
	EQ 10MG BASE	A077974 003	Jul 09, 2007
SUN PHARM INDUSTRIES	EQ 2.5MG BASE	A078081 001	Jan 31, 2008
	EQ 5MG BASE	A078081 002	Jan 31, 2008
	EQ 10MG BASE	A078081 003	Jan 31, 2008
SUNSHINE	EQ 2.5MG BASE	A206524 001	May 04, 2018
	EQ 5MG BASE	A206524 002	May 04, 2018
	EQ 10MG BASE	A206524 003	May 04, 2018
SYNTHON PHARMS	EQ 2.5MG BASE	A077080 001	Jun 27, 2007
	EQ 5MG BASE	A077080 002	Jun 27, 2007
	EQ 10MG BASE	A077080 003	Jun 27, 2007
TORRENT PHARMS	EQ 2.5MG BASE	A078573 001	Sep 22, 2008
	EQ 5MG BASE	A078573 002	Sep 22, 2008
	EQ 10MG BASE	A078573 003	Sep 22, 2008
WATSON LABS	EQ 2.5MG BASE	A077671 001	Jul 19, 2007
	EQ 5MG BASE	A077671 002	Jul 19, 2007
	EQ 10MG BASE	A077671 003	Jul 19, 2007
WOCKHARDT	EQ 2.5MG BASE	A078500 001	Sep 06, 2007
	EQ 5MG BASE	A078500 002	Sep 06, 2007
	EQ 10MG BASE	A078500 003	Sep 06, 2007

TABLET, ORALLY DISINTEGRATING;ORAL

AMLODIPINE BESYLATE

+ SYNTHON PHARMS	EQ 2.5MG BASE	N022026 001	Sep 27, 2007
+	EQ 5MG BASE	N022026 002	Sep 27, 2007
+	EQ 10MG BASE	N022026 003	Sep 27, 2007

AMLODIPINE BESYLATE; ATORVASTATIN CALCIUM

TABLET;ORAL

AMLODIPINE BESYLATE AND ATORVASTATIN CALCIUM

MYLAN	EQ 2.5MG BASE;EQ 10MG BASE	A200465 001	Nov 29, 2013
	EQ 2.5MG BASE;EQ 20MG BASE	A200465 002	Nov 29, 2013
	EQ 2.5MG BASE;EQ 40MG BASE	A200465 003	Nov 29, 2013

AMLODIPINE BESYLATE; BENAZEPRIL HYDROCHLORIDE

CAPSULE;ORAL

AMLODIPINE BESYLATE AND BENAZEPRIL HYDROCHLORIDE

AMNEAL	EQ 2.5MG BASE;10MG	A077215 001	Dec 07, 2018
	EQ 5MG BASE;10MG	A077215 002	Dec 07, 2018
	EQ 5MG BASE;20MG	A077215 003	Dec 07, 2018
	EQ 10MG BASE;20MG	A077215 004	Dec 07, 2018
NORVIUM BIOSCIENCE	EQ 2.5MG BASE;10MG	A077375 001	May 21, 2010
	EQ 5MG BASE;10MG	A077375 002	May 21, 2010
	EQ 5MG BASE;20MG	A077375 003	May 21, 2010
	EQ 5MG BASE;40MG	A079047 001	Jul 05, 2011
	EQ 10MG BASE;20MG	A077375 004	May 21, 2010

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

AMLODIPINE BESYLATE; BENAZEPRIL HYDROCHLORIDE

CAPSULE;ORAL

AMLODIPINE BESYLATE AND BENAZEPRIL HYDROCHLORIDE

	EQ 10MG BASE;40MG	A079047 002	Jul 05, 2011
STRIDES PHARMA	EQ 2.5MG BASE;10MG	A078381 001	Jul 29, 2010
	EQ 5MG BASE;10MG	A078381 002	Jul 29, 2010
	EQ 5MG BASE;20MG	A078381 003	Jul 29, 2010
	EQ 5MG BASE;40MG	A078381 005	Jul 29, 2010
	EQ 10MG BASE;20MG	A078381 004	Jul 29, 2010
	EQ 10MG BASE;40MG	A078381 006	Jul 29, 2010
TEVA PHARMS	EQ 2.5MG BASE;10MG	A077179 001	May 18, 2007
	EQ 5MG BASE;10MG	A077179 002	May 18, 2007
	EQ 5MG BASE;20MG	A077179 003	May 18, 2007
	EQ 5MG BASE;40MG	A077179 005	Jul 05, 2011
	EQ 10MG BASE;20MG	A077179 004	May 18, 2007
	EQ 10MG BASE;40MG	A077179 006	Jul 05, 2011

AMLODIPINE BESYLATE; CELECOXIB

TABLET;ORAL

CONSENSI

+	PURPLE BIOTECH	EQ 2.5MG BASE;200MG	N210045 001	May 31, 2018
+		EQ 5MG BASE;200MG	N210045 002	May 31, 2018
+		EQ 10MG BASE;200MG	N210045 003	May 31, 2018

AMLODIPINE BESYLATE; HYDROCHLOROTHIAZIDE; VALSARTAN

TABLET;ORAL

AMLODIPINE BESYLATE, VALSARTAN AND HYDROCHLOROTHIAZIDE

TEVA PHARMS	EQ 5MG BASE;12.5MG;160MG	A200435 001	Sep 25, 2012
	EQ 5MG BASE;25MG;160MG	A200435 002	Sep 25, 2012
	EQ 10MG BASE;12.5MG;160MG	A200435 005	Sep 25, 2012
	EQ 10MG BASE;25MG;160MG	A200435 003	Sep 25, 2012
	EQ 10MG BASE;25MG;320MG	A200435 004	Sep 25, 2012
TORRENT	EQ 5MG BASE;12.5MG;160MG	A201593 001	Jun 03, 2015
	EQ 5MG BASE;25MG;160MG	A201593 002	Jun 03, 2015
	EQ 10MG BASE;12.5MG;160MG	A201593 003	Jun 03, 2015
	EQ 10MG BASE;25MG;160MG	A201593 004	Jun 03, 2015
	EQ 10MG BASE;25MG;320MG	A201593 005	Jun 03, 2015

AMLODIPINE BESYLATE; OLMESARTAN MEDOXOMIL

TABLET;ORAL

AMLODIPINE AND OLMESARTAN MEDOXOMIL

ACCORD HLTHCARE INC	EQ 5MG BASE;20MG	A209600 001	Aug 30, 2018
	EQ 5MG BASE;40MG	A209600 003	Aug 30, 2018
	EQ 10MG BASE;20MG	A209600 002	Aug 30, 2018
	EQ 10MG BASE;40MG	A209600 004	Aug 30, 2018
GLENMARK PHARMS LTD	EQ 5MG BASE;20MG	A207807 001	Jul 05, 2017
	EQ 5MG BASE;40MG	A207807 002	Jul 05, 2017
	EQ 10MG BASE;20MG	A207807 003	Jul 05, 2017
	EQ 10MG BASE;40MG	A207807 004	Jul 05, 2017
JUBILANT GENERICS	EQ 5MG BASE;20MG	A207450 001	May 15, 2017
	EQ 5MG BASE;40MG	A207450 002	May 15, 2017
	EQ 10MG BASE;20MG	A207450 003	May 15, 2017
	EQ 10MG BASE;40MG	A207450 004	May 15, 2017
TEVA PHARMS USA	EQ 5MG BASE;20MG	A091154 001	Oct 26, 2016
	EQ 5MG BASE;40MG	A091154 002	Oct 26, 2016
	EQ 10MG BASE;20MG	A091154 003	Oct 26, 2016
	EQ 10MG BASE;40MG	A091154 004	Oct 26, 2016
TORRENT	EQ 5MG BASE;20MG	A202933 001	Nov 25, 2016
	EQ 5MG BASE;40MG	A202933 002	Nov 25, 2016
	EQ 10MG BASE;20MG	A202933 003	Nov 25, 2016
	EQ 10MG BASE;40MG	A202933 004	Nov 25, 2016

AMLODIPINE BESYLATE; PERINDOPRIL ARGININE

TABLET;ORAL

PRESTALIA

+	ADHERA	EQ 2.5MG BASE;3.5MG	N205003 001	Jan 21, 2015
+		EQ 5MG BASE;7MG	N205003 002	Jan 21, 2015
+		EQ 10MG BASE;14MG	N205003 003	Jan 21, 2015

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

AMLODIPINE BESYLATE; TELMISARTAN

TABLET; ORAL

TELMISARTAN AND AMLODIPINE

ALEMBIC	EQ 5MG BASE;40MG	A205234 001	Nov 17, 2016
	EQ 5MG BASE;80MG	A205234 003	Nov 17, 2016
	EQ 10MG BASE;40MG	A205234 002	Nov 17, 2016
	EQ 10MG BASE;80MG	A205234 004	Nov 17, 2016
TORRENT	EQ 5MG BASE;40MG	A202517 001	Jan 08, 2014
	EQ 5MG BASE;80MG	A202517 003	Jan 08, 2014
	EQ 10MG BASE;40MG	A202517 002	Jan 08, 2014
	EQ 10MG BASE;80MG	A202517 004	Jan 08, 2014
TWYNSTA			
+	BOEHRINGER INGELHEIM EQ 5MG BASE;40MG **	N022401 001	Oct 16, 2009
+	EQ 5MG BASE;80MG **	N022401 003	Oct 16, 2009
+	EQ 10MG BASE;40MG **	N022401 002	Oct 16, 2009
+	EQ 10MG BASE;80MG **	N022401 004	Oct 16, 2009

AMLODIPINE BESYLATE; VALSARTAN

TABLET; ORAL

AMLODIPINE BESYLATE AND VALSARTAN

TEVA PHARMS USA	EQ 5MG BASE;160MG	A091235 001	Mar 30, 2015
	EQ 5MG BASE;320MG	A091235 003	Mar 30, 2015
	EQ 10MG BASE;160MG	A091235 002	Mar 30, 2015
	EQ 10MG BASE;320MG	A091235 004	Mar 30, 2015
TORRENT	EQ 5MG BASE;160MG	A202377 001	Mar 30, 2015
	EQ 5MG BASE;320MG	A202377 002	Mar 30, 2015
	EQ 10MG BASE;160MG	A202377 003	Mar 30, 2015
	EQ 10MG BASE;320MG	A202377 004	Mar 30, 2015

AMLODIPINE MALEATE

TABLET; ORAL

AMVAZ

DR REDDYS LABS INC	2.5MG	N021435 001	Oct 31, 2003
	5MG	N021435 002	Oct 31, 2003
	10MG	N021435 003	Oct 31, 2003

AMMONIA N-13

INJECTABLE; INTRAVENOUS

AMMONIA N 13

CENTRAL RADIOPHARM	3.75-260mCi/ML	A204539 001	Jun 23, 2015
ESSENTIAL ISOTOPES	3.75mCi-260mCi/ML	A205687 001	Dec 17, 2015
SHERTECH LABS LLC	3.75mCi-260mCi/ML	A204366 001	Sep 19, 2014
SOFIE	18.8mCi-188mCi/5ML (3.75-37.5mCi/ML)	A204667 001	Apr 22, 2015
UNIV TX MD ANDERSON	30mCi-300mCi/8ML (3.75-37.5mCi/ML)	A203933 001	Jun 27, 2014
WISCONSIN	3.75mCi-260mCi/ML	A204356 001	Dec 18, 2014

AMMONIUM CHLORIDE

INJECTABLE; INJECTION

AMMONIUM CHLORIDE

ABBOTT	5MEQ/ML	A083130 001	
GD SEARLE LLC	3MEQ/ML	A086205 001	
AMMONIUM CHLORIDE 0.9% IN NORMAL SALINE			
MCGAW	900MG/100ML	N006580 001	
AMMONIUM CHLORIDE 2.14%			
B BRAUN	40MEQ/100ML	A085734 001	

AMMONIUM LACTATE

CREAM; TOPICAL

AMMONIUM LACTATE

WATSON LABS INC	EQ 12% BASE	A076829 001	Feb 07, 2006
LAC-HYDRIN			
+	SUN PHARM INDS INC EQ 12% BASE **	N020508 001	Aug 29, 1996
LOTION; TOPICAL			
AMMONIUM LACTATE			
WATSON LABS INC	EQ 12% BASE	A075575 001	Jun 11, 2002
LAC-HYDRIN			
+	SUN PHARM INDS INC EQ 12% BASE **	N019155 001	Apr 24, 1985

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

AMODIAQUINE HYDROCHLORIDE

TABLET;ORAL

CAMOQUIN HYDROCHLORIDE

PARKE DAVIS

EQ 200MG BASE

N006441 001

AMOXAPINE

TABLET;ORAL

AMOXAPINE

WATSON PHARMS TEVA

25MG
50MG
100MG
150MG

A072418 001 Aug 01, 1989
A072419 001 Aug 01, 1989
A072420 001 May 11, 1989
A072421 001 May 11, 1989

ASENDIN

LEDERLE

25MG
50MG
100MG
150MG

N018021 001
N018021 002
N018021 003
N018021 004

AMOXICILLIN

CAPSULE;ORAL

AMOXICILLIN

LABS ATRAL

250MG
500MG

A062528 001 Aug 07, 1985
A062528 002 Aug 07, 1985
A062067 001
A062067 002

MYLAN

250MG
500MG

A062884 001 Feb 25, 1988
A062881 001 Feb 25, 1988

STRIDES PHARMA

250MG
500MG

A065016 001 Apr 08, 1999
A065016 002 Apr 08, 1999

SUN PHARM INDS LTD

250MG
500MG

A062853 001 Dec 22, 1987
A063030 001 Feb 28, 1989

TEVA

250MG
500MG
500MG

A062854 001 Dec 22, 1987
A063031 001 Feb 28, 1989

AMOXIL

+ GLAXOSMITHKLINE

250MG **

N050459 001

+

500MG **

N050459 002

TRIMOX

APOTHECON

250MG
250MG
250MG
250MG
500MG
500MG
500MG
500MG

A061885 001
A062098 001
A062152 001
A063099 001 Mar 20, 1992
A061885 002
A062098 002
A062152 002
A063099 002 Mar 20, 1992

UTIMOX

PARKE DAVIS

250MG
500MG

A062107 001
A062107 002

WYMOX

WYETH AYERST

250MG
500MG

A062120 001
A062120 002

FOR SUSPENSION;ORAL

AMOXICILLIN

CHARTWELL RX

125MG/5ML
250MG/5ML

A062059 001
A062059 002

ENDO OPERATIONS

125MG/5ML
250MG/5ML

A062927 001 Nov 25, 1988
A062927 002 Nov 25, 1988

MYLAN

125MG/5ML
250MG/5ML

A062090 001
A062090 002

SUN PHARM INDS LTD

200MG/5ML
400MG/5ML

A065113 001 Nov 29, 2002
A065113 002 Nov 29, 2002

TEVA

125MG/5ML
125MG/5ML
250MG/5ML

A061931 001
A062946 001 Nov 01, 1988
A063001 001 Jan 06, 1989

AMOXIL

+ GLAXOSMITHKLINE

50MG/ML **

N050460 005

+

125MG/5ML **

N050460 001

+

250MG/5ML **

N050460 002

+

US ANTIBIOTICS

200MG/5ML **

N050760 001 Apr 15, 1999

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

AMOXICILLIN

FOR SUSPENSION;ORAL

LAROTID					
+	GLAXOSMITHKLINE	50MG/ML **		N050460 006	
POLYMOX					
	APOTHECON	125MG/5ML		A061851 001	
		125MG/5ML		A062323 001	
		250MG/5ML		A061851 002	
		250MG/5ML		A062323 002	
TRIMOX					
	APOTHECON	50MG/ML		A061886 001	
		125MG/5ML		A061886 002	
		125MG/5ML		A062099 001	
		125MG/5ML		A062154 001	
		125MG/5ML		A062885 001	Mar 08, 1988
		250MG/5ML		A061886 003	
		250MG/5ML		A062099 002	
		250MG/5ML		A062154 002	
		250MG/5ML		A062885 002	Mar 08, 1988
UTIMOX					
	PARKE DAVIS	125MG/5ML		A062127 001	
		250MG/5ML		A062127 002	
WYMOX					
	WYETH AYERST	125MG/5ML		A062131 001	
		250MG/5ML		A062131 002	

TABLET;ORAL

AMOXICILLIN					
	CHARTWELL RX	875MG		A065344 001	Jan 15, 2009
	SUN PHARM INDS LTD	500MG		A065059 001	Nov 24, 2000
		875MG		A065059 002	Nov 24, 2000
AMOXIL					
+	US ANTIBIOTICS	500MG **		N050754 002	Jul 10, 1998
+		875MG **		N050754 001	Jul 10, 1998
TABLET, CHEWABLE;ORAL					
AMOXICILLIN					
	APOTHECON	125MG		A064131 001	May 06, 1996
		250MG		A064131 002	May 06, 1996
	CHARTWELL RX	125MG		A064139 001	Jan 29, 1996
		250MG		A064139 002	Jan 29, 1996
	HIKMA	125MG		A205274 001	Jun 25, 2020
		250MG		A205274 002	Jun 25, 2020
	SUN PHARM INDS LTD	125MG		A065021 001	Dec 23, 1999
		200MG		A065060 001	Nov 29, 2000
		250MG		A065021 002	Dec 23, 1999
		400MG		A065060 002	Nov 29, 2000
	TEVA	125MG		A064031 001	Dec 19, 1996
		250MG		A064031 002	Dec 19, 1996
AMOXIL					
+	US ANTIBIOTICS	125MG **		N050542 002	
		200MG		N050761 001	Apr 15, 1999
+		250MG **		N050542 001	
		400MG		N050761 002	Apr 15, 1999
TABLET, EXTENDED RELEASE;ORAL					
MOXATAG					
+	PRAGMA	775MG		N050813 001	Jan 23, 2008
TABLET, FOR SUSPENSION;ORAL					
AMOXICILLIN					
	AUROBINDO PHARMA LTD	200MG		A065324 001	Jan 17, 2007
		400MG		A065324 002	Jan 17, 2007
DISPERMOX					
	RANBAXY LABS LTD	200MG		A065080 002	Aug 11, 2003
		400MG		A065080 001	Aug 11, 2003
		600MG		A065159 001	Dec 04, 2003

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

AMOXICILLIN; CLARITHROMYCIN; LANSOPRAZOLE

CAPSULE, TABLET, CAPSULE, DELAYED REL PELLETS;ORAL

LANSOPRAZOLE, AMOXICILLIN AND CLARITHROMYCIN (COPACKAGED)

ANI PHARMS 500MG;500MG;30MG

A200218 001 Aug 30, 2013

PREVPAC (COPACKAGED)

+ TAKEDA PHARMS USA 500MG;500MG;30MG **

N050757 001 Dec 02, 1997

AMOXICILLIN; CLARITHROMYCIN; OMEPRAZOLE

CAPSULE, TABLET, CAPSULE, DELAYED RELEASE;ORAL

OMECLAMOX-PAK

+ CUMBERLAND 500MG,N/A,N/A;N/A,500MG,N/A;N/A,N/A,20MG **

N050824 001 Feb 08, 2011

AMOXICILLIN; CLAVULANATE POTASSIUM

FOR SUSPENSION;ORAL

AMOXICILLIN AND CLAVULANATE POTASSIUM

SUN PHARM INDS LTD 200MG/5ML;EQ 28.5MG BASE/5ML

A065132 001 Mar 19, 2003

400MG/5ML;EQ 57MG BASE/5ML

A065132 002 Mar 19, 2003

600MG/5ML;EQ 42.9MG BASE/5ML

A065207 002 Jan 30, 2007

AUGMENTIN '200'

+ US ANTIBIOTICS 200MG/5ML;EQ 28.5MG BASE/5ML

N050725 001 May 31, 1996

AUGMENTIN '400'

+ US ANTIBIOTICS 400MG/5ML;EQ 57MG BASE/5ML

N050725 002 May 31, 1996

TABLET;ORAL

AMOXICILLIN AND CLAVULANATE POTASSIUM

APOTEX INC 250MG;EQ 125MG BASE

A065333 001 Feb 24, 2009

500MG;EQ 125MG BASE

A065333 002 Feb 24, 2009

875MG;EQ 125MG BASE

A065317 003 Oct 20, 2008

SANDOZ INC 500MG;EQ 125MG BASE

A065117 001 Nov 27, 2002

SUN PHARM INDS LTD 500MG;EQ 125MG BASE

A065109 001 Nov 04, 2002

875MG;EQ 125MG BASE

A065102 001 Sep 17, 2002

AUGMENTIN '250'

+ US ANTIBIOTICS 250MG;EQ 125MG BASE **

N050564 001 Aug 06, 1984

AUGMENTIN '500'

+ US ANTIBIOTICS 500MG;EQ 125MG BASE **

N050564 002 Aug 06, 1984

TABLET, CHEWABLE;ORAL

AMOXICILLIN AND CLAVULANATE POTASSIUM

CHARTWELL RX 200MG;EQ 28.5MG BASE

A065065 001 Apr 18, 2002

400MG;EQ 57MG BASE

A065065 002 Apr 18, 2002

SUN PHARM INDS LTD 200MG;EQ 28.5MG BASE

A065161 001 Dec 03, 2003

400MG;EQ 57MG BASE

A065161 002 Dec 03, 2003

AUGMENTIN '125'

+ US ANTIBIOTICS 125MG;EQ 31.25MG BASE **

N050597 001 Jul 22, 1985

AUGMENTIN '200'

+ US ANTIBIOTICS 200MG;EQ 28.5MG BASE

N050726 001 May 31, 1996

AUGMENTIN '250'

+ US ANTIBIOTICS 250MG;EQ 62.5MG BASE **

N050597 002 Jul 22, 1985

AUGMENTIN '400'

+ US ANTIBIOTICS 400MG;EQ 57MG BASE

N050726 002 May 31, 1996

TABLET, EXTENDED RELEASE;ORAL

AUGMENTIN XR

+ US ANTIBIOTICS 1GM;EQ 62.5MG BASE **

N050785 001 Sep 25, 2002

AMPHETAMINE

SUSPENSION, EXTENDED RELEASE;ORAL

ADZENYS ER

+ NEOS THERAPS INC EQ 1.25MG BASE/ML

N204325 001 Sep 15, 2017

TABLET, ORALLY DISINTEGRATING, EXTENDED RELEASE;ORAL

AMPHETAMINE

ACTAVIS LABS FL INC EQ 3.1MG BASE

A209253 001 Jun 22, 2023

EQ 6.3MG BASE

A209253 002 Jun 22, 2023

EQ 9.4MG BASE

A209253 003 Jun 22, 2023

EQ 12.5MG BASE

A209253 004 Jun 22, 2023

EQ 15.7MG BASE

A209253 005 Jun 22, 2023

EQ 18.8MG BASE

A209253 006 Jun 22, 2023

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

AMPHETAMINE ADIPATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE ADIPATE; DEXTROAMPHETAMINE SULFATE

CAPSULE; ORAL

DELCOBESE

TEVA	1.25MG; 1.25MG; 1.25MG; 1.25MG **	A083564	001
	2.5MG; 2.5MG; 2.5MG; 2.5MG **	A083564	002
	3.75MG; 3.75MG; 3.75MG; 3.75MG **	A083564	003
	5MG; 5MG; 5MG; 5MG **	A083564	004

TABLET; ORAL

DELCOBESE

TEVA	1.25MG; 1.25MG; 1.25MG; 1.25MG	A083563	004
	2.5MG; 2.5MG; 2.5MG; 2.5MG	A083563	003
	3.75MG; 3.75MG; 3.75MG; 3.75MG	A083563	002
	5MG; 5MG; 5MG; 5MG	A083563	001

AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATE

CAPSULE, EXTENDED RELEASE; ORAL

DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE

AUROLIFE PHARMA LLC	1.25MG; 1.25MG; 1.25MG; 1.25MG	A211876	001	Jun 24, 2020
	2.5MG; 2.5MG; 2.5MG; 2.5MG	A211876	002	Jun 24, 2020
	3.75MG; 3.75MG; 3.75MG; 3.75MG	A211876	003	Jun 24, 2020
	5MG; 5MG; 5MG; 5MG	A211876	004	Jun 24, 2020
	6.25MG; 6.25MG; 6.25MG; 6.25MG	A211876	005	Jun 24, 2020
	7.5MG; 7.5MG; 7.5MG; 7.5MG	A211876	006	Jun 24, 2020
ENDO OPERATIONS	1.25MG; 1.25MG; 1.25MG; 1.25MG	A206159	001	May 31, 2019
	2.5MG; 2.5MG; 2.5MG; 2.5MG	A206159	002	May 31, 2019
	3.75MG; 3.75MG; 3.75MG; 3.75MG	A206159	003	May 31, 2019
	5MG; 5MG; 5MG; 5MG	A206159	004	May 31, 2019
	6.25MG; 6.25MG; 6.25MG; 6.25MG	A206159	005	May 31, 2019
	7.5MG; 7.5MG; 7.5MG; 7.5MG	A206159	006	May 31, 2019
NESHER PHARMS	1.25MG; 1.25MG; 1.25MG; 1.25MG	A210080	001	Jul 17, 2019
	2.5MG; 2.5MG; 2.5MG; 2.5MG	A210080	002	Jul 17, 2019
	3.75MG; 3.75MG; 3.75MG; 3.75MG	A210080	003	Jul 17, 2019
	5MG; 5MG; 5MG; 5MG	A210080	004	Jul 17, 2019
	6.25MG; 6.25MG; 6.25MG; 6.25MG	A210080	005	Jul 17, 2019
	7.5MG; 7.5MG; 7.5MG; 7.5MG	A210080	006	Jul 17, 2019
SUN PHARM INDUSTRIES	1.25MG; 1.25MG; 1.25MG; 1.25MG	A211715	001	May 17, 2019
	2.5MG; 2.5MG; 2.5MG; 2.5MG	A211715	002	May 17, 2019
	3.75MG; 3.75MG; 3.75MG; 3.75MG	A211715	003	May 17, 2019
	5MG; 5MG; 5MG; 5MG	A211715	004	May 17, 2019
	6.25MG; 6.25MG; 6.25MG; 6.25MG	A211715	005	May 17, 2019
	7.5MG; 7.5MG; 7.5MG; 7.5MG	A211715	006	May 17, 2019
TEVA	1.25MG; 1.25MG; 1.25MG; 1.25MG	A077488	001	Apr 29, 2013
	2.5MG; 2.5MG; 2.5MG; 2.5MG	A077488	002	Apr 29, 2013
	3.75MG; 3.75MG; 3.75MG; 3.75MG	A077488	003	Apr 29, 2013
	5MG; 5MG; 5MG; 5MG	A077488	004	Apr 29, 2013
	6.25MG; 6.25MG; 6.25MG; 6.25MG	A077488	005	Apr 29, 2013
	7.5MG; 7.5MG; 7.5MG; 7.5MG	A077488	006	Apr 29, 2013
DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE				
BARR LABS INC	1.25MG; 1.25MG; 1.25MG; 1.25MG	A076536	001	Feb 12, 2013
	2.5MG; 2.5MG; 2.5MG; 2.5MG	A076536	002	Feb 12, 2013
	3.75MG; 3.75MG; 3.75MG; 3.75MG	A076536	003	Feb 12, 2013
	5MG; 5MG; 5MG; 5MG	A076536	004	Feb 12, 2013
	6.25MG; 6.25MG; 6.25MG; 6.25MG	A076536	005	Feb 12, 2013
	7.5MG; 7.5MG; 7.5MG; 7.5MG	A076536	006	Feb 12, 2013

TABLET; ORAL

ADDERALL 10

+ TEVA WOMENS 2.5MG; 2.5MG; 2.5MG; 2.5MG ** N011522 007 Feb 13, 1996

ADDERALL 12.5

+ TEVA WOMENS 3.125MG; 3.125MG; 3.125MG; 3.125MG ** N011522 012 Aug 31, 2000

ADDERALL 15

+ TEVA WOMENS 3.75MG; 3.75MG; 3.75MG; 3.75MG ** N011522 013 Aug 31, 2000

ADDERALL 20

+ TEVA WOMENS 5MG; 5MG; 5MG; 5MG ** N011522 008 Feb 13, 1996

ADDERALL 30

+ TEVA WOMENS 7.5MG; 7.5MG; 7.5MG; 7.5MG ** N011522 010 May 12, 1997

ADDERALL 5

+ TEVA WOMENS 1.25MG; 1.25MG; 1.25MG; 1.25MG ** N011522 009 May 12, 1997

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATE

TABLET; ORAL

ADDERALL 7.5

+ TEVA WOMENS 1.875MG;1.875MG;1.875MG;1.875MG ** N011522 011 Aug 31, 2000

DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE

ACCORD HLTHCARE 1.25MG;1.25MG;1.25MG;1.25MG A214347 001 Nov 22, 2021

1.875MG;1.875MG;1.875MG;1.875MG A214347 002 Nov 22, 2021

2.5MG;2.5MG;2.5MG;2.5MG A214347 003 Nov 22, 2021

3.125MG;3.125MG;3.125MG;3.125MG A214347 004 Nov 22, 2021

3.75MG;3.75MG;3.75MG;3.75MG A214347 005 Nov 22, 2021

5MG;5MG;5MG;5MG A214347 006 Nov 22, 2021

7.5MG;7.5MG;7.5MG;7.5MG A214347 007 Nov 22, 2021

ACTAVIS ELIZABETH 1.25MG;1.25MG;1.25MG;1.25MG A040456 001 May 06, 2003

2.5MG;2.5MG;2.5MG;2.5MG A040456 002 May 06, 2003

5MG;5MG;5MG;5MG A040456 003 May 06, 2003

7.5MG;7.5MG;7.5MG;7.5MG A040456 004 May 06, 2003

CEDIPROF INC 1.25MG;1.25MG;1.25MG;1.25MG A210754 001 Jul 05, 2022

2.5MG;2.5MG;2.5MG;2.5MG A210754 002 Jul 05, 2022

3.75MG;3.75MG;3.75MG;3.75MG A210754 003 Jul 05, 2022

5MG;5MG;5MG;5MG A210754 004 Jul 05, 2022

7.5MG;7.5MG;7.5MG;7.5MG A210754 005 Jul 05, 2022

NORVIUM BIOSCIENCE 1.25MG;1.25MG;1.25MG;1.25MG A206721 001 Nov 10, 2015

1.875MG;1.875MG;1.875MG;1.875MG A206721 002 Nov 10, 2015

2.5MG;2.5MG;2.5MG;2.5MG A206721 003 Nov 10, 2015

3.125MG;3.125MG;3.125MG;3.125MG A206721 004 Nov 10, 2015

3.75MG;3.75MG;3.75MG;3.75MG A206721 005 Nov 10, 2015

5MG;5MG;5MG;5MG A206721 006 Nov 10, 2015

7.5MG;7.5MG;7.5MG;7.5MG A206721 007 Nov 10, 2015

TEVA PHARMS 1.25MG;1.25MG;1.25MG;1.25MG A040472 001 Sep 30, 2003

2.5MG;2.5MG;2.5MG;2.5MG A040472 002 Sep 30, 2003

5MG;5MG;5MG;5MG A040472 003 Sep 30, 2003

7.5MG;7.5MG;7.5MG;7.5MG A040472 004 Sep 30, 2003

AMPHETAMINE RESIN COMPLEX; DEXTROAMPHETAMINE RESIN COMPLEX

CAPSULE, EXTENDED RELEASE; ORAL

BIPHETAMINE 12.5

UCB INC EQ 6.25MG BASE;EQ 6.25MG BASE N010093 007

BIPHETAMINE 20

UCB INC EQ 10MG BASE;EQ 10MG BASE N010093 003

BIPHETAMINE 7.5

UCB INC EQ 3.75MG BASE;EQ 3.75MG BASE N010093 009

AMPHETAMINE SULFATE

TABLET; ORAL

AMPHETAMINE SULFATE

DR REDDYS LABS SA 5MG A213898 001 Jul 14, 2020

10MG A213898 002 Jul 14, 2020

GLENMARK PHARMS LTD 5MG A212186 001 Jan 27, 2021

10MG A212186 002 Jan 27, 2021

+ LANNETT 5MG A083901 001 Aug 31, 1984

+ 10MG A083901 002 Aug 31, 1984

NOVAST LABS 5MG A213763 001 Aug 24, 2020

10MG A213763 002 Aug 24, 2020

SUN PHARM INDS INC 5MG A214574 001 Jan 27, 2021

10MG A214574 002 Jan 27, 2021

TABLET, ORALLY DISINTEGRATING; ORAL

EVEKEO ODT

+ AZURITY 2.5MG ** N209905 005 Apr 16, 2021

+ 5MG N209905 001 Jan 30, 2019

+ 10MG N209905 002 Jan 30, 2019

+ 15MG N209905 003 Jan 30, 2019

+ 20MG N209905 004 Jan 30, 2019

AMPHOTERICIN B

CREAM; TOPICAL

FUNGIZONE

APOTHECON 3% N050314 001

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

AMPHOTERICIN B

INJECTABLE; INJECTION

AMPHOTERICIN B

ABBOTT	50MG/VIAL	A064141	001	Dec 23, 1996
ABRAXIS PHARM	50MG/VIAL	A062728	001	Apr 13, 1987
TEVA PARENTERAL	50MG/VIAL	A064062	001	Mar 31, 1995

FUNGIZONE

APOTHECON	50MG/VIAL	A060517	001	
-----------	-----------	---------	-----	--

INJECTABLE, LIPID COMPLEX; INJECTION

AMPHOTEC

ALKOPHARMA USA	50MG/VIAL	N050729	001	Nov 22, 1996
	100MG/VIAL	N050729	002	Nov 22, 1996

LOTION; TOPICAL

FUNGIZONE

APOTHECON	3%	A060570	001	
-----------	----	---------	-----	--

OINTMENT; TOPICAL

FUNGIZONE

APOTHECON	3%	N050313	001	
-----------	----	---------	-----	--

SUSPENSION; ORAL

FUNGIZONE

BRISTOL MYERS SQUIBB	100MG/ML	N050341	003	
----------------------	----------	---------	-----	--

AMPICILLIN SODIUM

INJECTABLE; INJECTION

AMPICILLIN SODIUM

ACS DOBFAR SPA	EQ 500MG BASE/VIAL	A090884	001	Apr 03, 2013
	EQ 1GM BASE/VIAL	A090884	002	Apr 03, 2013
	EQ 2GM BASE/VIAL	A090884	003	Apr 03, 2013
APOTHECON	EQ 125MG BASE/VIAL	A062860	001	Feb 05, 1988
	EQ 250MG BASE/VIAL	A062860	002	Feb 05, 1988
	EQ 500MG BASE/VIAL	A062860	003	Feb 05, 1988
	EQ 1GM BASE/VIAL	A062860	004	Feb 05, 1988
	EQ 2GM BASE/VIAL	A062860	005	Feb 05, 1988
CONSOLIDATED PHARM	EQ 125MG BASE/VIAL	A061936	005	
	EQ 250MG BASE/VIAL	A061936	001	
	EQ 500MG BASE/VIAL	A061936	002	
	EQ 1GM BASE/VIAL	A061936	003	
	EQ 2GM BASE/VIAL	A061936	004	
EUGIA PHARMA SPECLTS	EQ 125MG BASE/VIAL	A065499	001	Aug 17, 2010
HIKMA	EQ 125MG BASE/VIAL	A062692	001	Jun 24, 1986
	EQ 250MG BASE/VIAL	A062692	002	Jun 24, 1986
	EQ 500MG BASE/VIAL	A062692	003	Jun 24, 1986
	EQ 1GM BASE/VIAL	A062692	004	Jun 24, 1986
	EQ 2GM BASE/VIAL	A062692	005	Jun 24, 1986
HOSPIRA	EQ 10GM BASE/VIAL	A062692	006	Jun 24, 1986
	EQ 250MG BASE/VIAL	A202864	001	Sep 04, 2015
	EQ 500MG BASE/VIAL	A202864	002	Sep 04, 2015
	EQ 1GM BASE/VIAL	A202864	003	Sep 04, 2015
	EQ 2GM BASE/VIAL	A202864	004	Sep 04, 2015
HQ SPECLT PHARMA	EQ 10GM BASE/VIAL	A202865	001	Sep 04, 2015
	EQ 125MG BASE/VIAL	A062772	005	Apr 15, 1993
	EQ 500MG BASE/VIAL	A062772	008	Apr 15, 1993
	EQ 1GM BASE/VIAL	A062772	002	Apr 15, 1993
	EQ 2GM BASE/VIAL	A062772	004	Apr 15, 1993
INTL MEDICATION	EQ 1GM BASE/VIAL	A062634	002	Jan 09, 1987
	EQ 2GM BASE/VIAL	A062634	003	Jan 09, 1987
ISTITUTO BIO ITA SPA	EQ 125MG BASE/VIAL	A062797	001	Jul 12, 1993
LILLY	EQ 500MG BASE/VIAL	A062565	001	Apr 04, 1985
	EQ 1GM BASE/VIAL	A062565	002	Apr 04, 1985
	EQ 2GM BASE/VIAL	A062565	003	Jun 24, 1986
STERISCIENCE SPECLTS	EQ 250MG BASE/VIAL	A201025	001	Apr 09, 2014
	EQ 500MG BASE/VIAL	A201025	002	Apr 09, 2014
WATSON LABS INC	EQ 125MG BASE/VIAL	A062816	001	Oct 24, 1988
	EQ 250MG BASE/VIAL	A062816	002	Oct 24, 1988
	EQ 500MG BASE/VIAL	A062816	003	Oct 24, 1988
	EQ 1GM BASE/VIAL	A062816	004	Oct 24, 1988
	EQ 2GM BASE/VIAL	A062816	005	Oct 24, 1988
	EQ 10GM BASE/VIAL	A062994	001	Sep 15, 1988

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

AMPICILLIN SODIUMINJECTABLE; INJECTION
OMNIPEN-N

WYETH AYERST	EQ 125MG BASE/VIAL	A060626 001	
	EQ 125MG BASE/VIAL	A062718 001	Dec 16, 1986
	EQ 250MG BASE/VIAL	A060626 002	
	EQ 250MG BASE/VIAL	A062718 002	Dec 16, 1986
	EQ 500MG BASE/VIAL	A060626 003	
	EQ 500MG BASE/VIAL	A062718 003	Dec 16, 1986
	EQ 1GM BASE/VIAL	A060626 004	
	EQ 1GM BASE/VIAL	A062718 004	Dec 16, 1986
	EQ 2GM BASE/VIAL	A060626 005	
	EQ 2GM BASE/VIAL	A062718 005	Dec 16, 1986

PENBRITIN-S

+ WYETH AYERST	EQ 125MG BASE/VIAL **	N050072 001	
+	EQ 250MG BASE/VIAL **	N050072 002	
+	EQ 500MG BASE/VIAL **	N050072 003	
+	EQ 1GM BASE/VIAL **	N050072 004	
+	EQ 2GM BASE/VIAL **	N050072 005	
+	EQ 4GM BASE/VIAL **	N050072 006	

POLYCILLIN-N

BRISTOL	EQ 125MG BASE/VIAL **	N050309 001	
	EQ 250MG BASE/VIAL **	N050309 002	
	EQ 500MG BASE/VIAL **	N050309 003	
	EQ 1GM BASE/VIAL **	N050309 004	
	EQ 2GM BASE/VIAL **	N050309 005	

TOTACILLIN-N

GLAXOSMITHKLINE	EQ 125MG BASE/VIAL	A060677 001	
	EQ 250MG BASE/VIAL	A060677 002	
	EQ 500MG BASE/VIAL	A060677 003	
	EQ 1GM BASE/VIAL	A060677 004	
	EQ 1GM BASE/VIAL	A062727 001	Dec 19, 1986
	EQ 2GM BASE/VIAL	A060677 005	
	EQ 2GM BASE/VIAL	A062727 002	Dec 19, 1986
	EQ 10GM BASE/VIAL	A060677 006	

AMPICILLIN SODIUM; SULBACTAM SODIUM

INJECTABLE; INJECTION

AMPICILLIN AND SULBACTAM

ASTRAL	EQ 1GM BASE/VIAL;EQ 500MG BASE/VIAL	A090579 001	Jan 08, 2016
	EQ 2GM BASE/VIAL;EQ 1GM BASE/VIAL	A090579 002	Jan 08, 2016
	EQ 10GM BASE/VIAL;EQ 5GM BASE/VIAL	A090578 001	Jan 11, 2016
EUGIA PHARMA SPECLTS	EQ 1GM BASE/VIAL;EQ 500MG BASE/VIAL	A090340 001	Sep 20, 2010
	EQ 2GM BASE/VIAL;EQ 1GM BASE/VIAL	A090340 002	Sep 20, 2010
HOSPIRA INC	EQ 1GM BASE/VIAL;EQ 500MG BASE/VIAL	A090375 001	Dec 21, 2011
	EQ 1GM BASE/VIAL;EQ 500MG BASE/VIAL	A090653 001	Dec 21, 2011
	EQ 2GM BASE/VIAL;EQ 1GM BASE/VIAL	A090375 002	Dec 21, 2011
	EQ 2GM BASE/VIAL;EQ 1GM BASE/VIAL	A090653 002	Dec 21, 2011
	EQ 10GM BASE/VIAL;EQ 5GM BASE/VIAL	A090646 001	Dec 21, 2011
MEDIMETRIKS PHARMS	EQ 1GM BASE/VIAL;EQ 500MG BASE/VIAL	A065316 001	Jun 29, 2007
	EQ 2GM BASE/VIAL;EQ 1GM BASE/VIAL	A065316 002	Jun 29, 2007
UNASYN			
PFIZER	EQ 2GM BASE/VIAL;EQ 1GM BASE/VIAL	A062901 002	Feb 27, 1992
	EQ 500MG BASE/VIAL;EQ 250MG BASE/VIAL	N050608 003	Dec 31, 1986
	EQ 1GM BASE/VIAL;EQ 500MG BASE/VIAL	A062901 001	Nov 23, 1988

AMPICILLIN/AMPICILLIN TRIHYDRATE

CAPSULE; ORAL

AMCILL

PARKE DAVIS	EQ 250MG BASE	A062041 001	
	EQ 500MG BASE	A062041 002	

AMPICILLIN TRIHYDRATE

BELCHER PHARMS	EQ 250MG BASE	A061602 001	
	EQ 500MG BASE	A061602 002	
CHARTWELL RX	EQ 250MG BASE	A061502 001	
	EQ 500MG BASE	A061502 002	
IVAX SUB TEVA PHARMS	EQ 250MG BASE	A060765 001	
	EQ 500MG BASE	A060765 002	
LEDERLE	EQ 250MG BASE	A062208 001	
	EQ 500MG BASE	A062208 002	

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

AMPICILLIN/AMPICILLIN TRIHYDRATE

CAPSULE;ORAL

AMPICILLIN TRIHYDRATE			
MYLAN	EQ 250MG BASE	A061755	001
	EQ 500MG BASE	A061755	002
PUREPAC PHARM	EQ 250MG BASE	A061853	001
	EQ 500MG BASE	A061853	002
SANDOZ	EQ 250MG BASE	A064082	001 Aug 29, 1995
STRIDES PHARMA	EQ 250MG BASE	A062883	001 Feb 25, 1988
	EQ 500MG BASE	A062882	001 Feb 25, 1988
VITARINE	EQ 250MG BASE	A061387	001
	EQ 500MG BASE	A061387	003
OMNIPEN (AMPICILLIN)			
WYETH AYERST	250MG	A060624	001
	500MG	A060624	002
PENBRITIN			
WYETH AYERST	EQ 250MG BASE	A060908	001
	EQ 500MG BASE	A060908	002
PFIZERPEN-A			
PFIZER	EQ 250MG BASE	A062050	001
	EQ 500MG BASE	A062050	002
POLYCILLIN			
BRISTOL	EQ 250MG BASE	N050310	001
	EQ 500MG BASE	N050310	002
PRINCIPEN			
APOTHECON	EQ 250MG BASE	A062888	001 Mar 04, 1988
	EQ 500MG BASE	A062888	002 Mar 04, 1988
BRISTOL MYERS SQUIBB	EQ 250MG BASE	A061392	001
	EQ 500MG BASE	A061392	002
PRINCIPEN '250'			
APOTHECON	EQ 250MG BASE	A062157	002
+	EQ 250MG BASE	N050056	001
PRINCIPEN '500'			
APOTHECON	EQ 500MG BASE	A062157	001
+	EQ 500MG BASE	N050056	002
TOTACILLIN			
GLAXOSMITHKLINE	EQ 250MG BASE	A060060	001
	EQ 250MG BASE	A062212	001
	EQ 500MG BASE	A060060	002
	EQ 500MG BASE	A062212	002
FOR SUSPENSION;ORAL			
AMCILL			
PARKE DAVIS	EQ 125MG BASE/5ML	A062030	001
	EQ 250MG BASE/5ML	A062030	002
AMPICILLIN TRIHYDRATE			
BELCHER PHARMS	EQ 125MG BASE/5ML	A061601	001
	EQ 250MG BASE/5ML	A061601	002
ENDO OPERATIONS	EQ 125MG BASE/5ML	A062982	001 Feb 10, 1989
	EQ 250MG BASE/5ML	A062982	002 Feb 10, 1989
MYLAN	EQ 125MG BASE/5ML	A061829	002
	EQ 250MG BASE/5ML	A061829	001
PUREPAC PHARM	EQ 125MG BASE/5ML	A061980	001
	EQ 250MG BASE/5ML	A061980	002
TEVA	EQ 125MG BASE/5ML	A061370	001
	EQ 250MG BASE/5ML	A061370	002
OMNIPEN (AMPICILLIN)			
WYETH AYERST	100MG/ML	A060625	001
	125MG/5ML	A060625	002
	250MG/5ML	A060625	003
	500MG/5ML	A060625	004
PENBRITIN			
WYETH AYERST	EQ 100MG BASE/ML	N050019	001
	EQ 125MG BASE/5ML	N050019	002
	EQ 250MG BASE/5ML	N050019	003
PFIZERPEN-A			
PFIZER	EQ 125MG BASE/5ML	A062049	001
	EQ 250MG BASE/5ML	A062049	002
POLYCILLIN			
APOTHECON	EQ 125MG BASE/5ML	A062297	001

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

AMPICILLIN/AMPICILLIN TRIHYDRATE

FOR SUSPENSION;ORAL

POLYCILLIN

	EQ 250MG BASE/5ML	A062297 002
BRISTOL	EQ 100MG BASE/ML	N050308 004
	EQ 125MG BASE/5ML	N050308 001
	EQ 250MG BASE/5ML	N050308 002
	EQ 500MG BASE/5ML	N050308 003
PRINCIPEN		
APOTHECON	EQ 100MG BASE/ML	A061394 001
	EQ 125MG BASE/5ML	A061394 002
	EQ 250MG BASE/5ML	A061394 003
PRINCIPEN '125'		
APOTHECON	EQ 125MG BASE/5ML	A060127 002
	EQ 125MG BASE/5ML	A062151 001
PRINCIPEN '250'		
APOTHECON	EQ 250MG BASE/5ML	A060127 001
	EQ 250MG BASE/5ML	A062151 002
TOTACILLIN		
GLAXOSMITHKLINE	EQ 125MG BASE/5ML	A060666 001
	EQ 125MG BASE/5ML	A062223 001
	EQ 250MG BASE/5ML	A060666 002
	EQ 250MG BASE/5ML	A062223 002

TABLET, CHEWABLE;ORAL

POLYCILLIN

BRISTOL EQ 125MG BASE N050093 001

AMPICILLIN/AMPICILLIN TRIHYDRATE; PROBENECID

CAPSULE;ORAL

PRINCIPEN W/ PROBENECID

APOTHECON	EQ 389MG BASE;111MG	A062150 001
	EQ 389MG BASE;111MG	N050488 001

FOR SUSPENSION;ORAL

POLYCILLIN-PRB

APOTHECON	EQ 3.5GM BASE/BOT;1GM/BOT	A061898 001
BRISTOL	EQ 3.5GM BASE/BOT;1GM/BOT	N050457 001

PROBAMPACIN

COSETTE EQ 3.5GM BASE/BOT;1GM/BOT A061741 001

AMPRENAVIR

CAPSULE;ORAL

AGENERASE

GLAXOSMITHKLINE	50MG	N021007 001	Apr 15, 1999
	150MG	N021007 002	Apr 15, 1999

SOLUTION;ORAL

AGENERASE

+ GLAXOSMITHKLINE 15MG/ML ** N021039 001 Apr 15, 1999

ANAGRELIDE HYDROCHLORIDE

CAPSULE;ORAL

AGRYLIN

+ TAKEDA PHARMS USA EQ 1MG BASE ** N020333 002 Mar 14, 1997

ANAGRELIDE HYDROCHLORIDE

BARR	EQ 0.5MG BASE	A076530 001	Apr 18, 2005
	EQ 1MG BASE	A076530 002	Apr 18, 2005
CHARTWELL RX	EQ 0.5MG BASE	A076683 001	Apr 18, 2005
	EQ 1MG BASE	A076683 002	Apr 18, 2005
ESJAY PHARMA	EQ 0.5MG BASE	A076811 001	Apr 18, 2005
	EQ 1MG BASE	A076811 002	Apr 18, 2005
RISING	EQ 0.5MG BASE	A077613 001	Jun 27, 2006
	EQ 1MG BASE	A077613 002	Jun 27, 2006
ROXANE	EQ 0.5MG BASE	A076489 001	Apr 18, 2005
	EQ 1MG BASE	A076489 002	Apr 18, 2005
WATSON LABS	EQ 0.5MG BASE	A076417 001	Apr 18, 2005
	EQ 1MG BASE	A076417 002	Apr 18, 2005

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ANASTROZOLE

TABLET; ORAL

ANASTROZOLE

APOTEX INC	1MG	A200654 001	May 11, 2012
CHARTWELL MOLECULAR	1MG	A091331 001	Jan 05, 2011
CHARTWELL RX	1MG	A090732 001	Jun 28, 2010
FRESENIUS KABI USA	1MG	A090088 001	Jun 28, 2010
HIKMA	1MG	A078485 001	Jun 28, 2010
IMPAX LABS INC	1MG	A091242 001	May 31, 2012
NORVIUM BIOSCIENCE	1MG	A091051 001	Jun 28, 2010
SANDOZ	1MG	A079007 001	Jun 28, 2010
SUN PHARM INDS LTD	1MG	A091177 001	Jul 15, 2011
SYNTHON PHARMS	1MG	A078322 001	Jun 28, 2010
WATSON LABS TEVA	1MG	A078984 001	Jun 28, 2010

ANGIOTENSIN II ACETATE

SOLUTION; INTRAVENOUS

GIAPREZA

+ LA JOLLA PHARMA	EQ 5MG BASE/2ML (EQ 2.5MG BASE/ML)	N209360 002	Dec 21, 2017
-------------------	------------------------------------	-------------	--------------

ANILERIDINE HYDROCHLORIDE

TABLET; ORAL

LERITINE

MERCK	EQ 25MG BASE	N010585 002	
-------	--------------	-------------	--

ANILERIDINE PHOSPHATE

INJECTABLE; INJECTION

LERITINE

MERCK	25MG/ML	N010520 003	
-------	---------	-------------	--

ANISINDIONE

TABLET; ORAL

MIRADON

SCHERING	50MG	N010909 003	
----------	------	-------------	--

ANISOTROPINE METHYLBROMIDE

TABLET; ORAL

ANISOTROPINE METHYLBROMIDE

WATSON LABS	50MG	A086046 001	
-------------	------	-------------	--

VALPIN 50

ENDO PHARMS	50MG	N013428 001	
-------------	------	-------------	--

ANTAZOLINE PHOSPHATE; NAPHAZOLINE HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

VASOCON-A

NOVARTIS	0.5%; 0.05%	N018746 002	Jul 11, 1994
----------	-------------	-------------	--------------

APIXABAN

TABLET; ORAL

APIXABAN

APOTEX	2.5MG	A210091 001	Feb 16, 2024
	5MG	A210091 002	Feb 16, 2024
AUROBINDO PHARMA LTD	2.5MG	A210026 001	May 26, 2023
	5MG	A210026 002	May 26, 2023
BIONPHARMA	2.5MG	A210152 001	Apr 08, 2020
	5MG	A210152 002	Apr 08, 2020
BRECKENRIDGE	2.5MG	A209845 001	Jul 26, 2021
	5MG	A209845 002	Jul 26, 2021
MICRO LABS	2.5MG	A210013 001	Dec 23, 2019
	5MG	A210013 002	Dec 23, 2019
MYLAN	2.5MG	A210128 001	Dec 23, 2019
	5MG	A210128 002	Dec 23, 2019
ZYDUS PHARMS	2.5MG	A210185 001	Feb 27, 2023
	5MG	A210185 002	Feb 27, 2023

APOMORPHINE HYDROCHLORIDE

FILM; SUBLINGUAL

KYNMOBI

+ SUMITOMO PHARMA AM	10MG	N210875 001	May 21, 2020
	15MG	N210875 002	May 21, 2020
	20MG	N210875 003	May 21, 2020
	25MG	N210875 004	May 21, 2020
	30MG	N210875 005	May 21, 2020

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

APOMORPHINE HYDROCHLORIDE

INJECTABLE;SUBCUTANEOUS

APOKYN

MDD US

20MG/2ML (10MG/ML)

N021264 001 Apr 20, 2004

APREMILAST

TABLET;ORAL

APREMILAST

AMNEAL

10MG

A211782 001 Jun 30, 2021

20MG

A211782 002 Jun 30, 2021

30MG

A211782 003 Jun 30, 2021

AUROBINDO PHARMA LTD

10MG

A211716 001 Jul 19, 2023

20MG

A211716 002 Jul 19, 2023

30MG

A211716 003 Jul 19, 2023

DR REDDYS

10MG

A211756 001 Jul 14, 2023

20MG

A211756 002 Jul 14, 2023

30MG

A211756 003 Jul 14, 2023

TEVA PHARMS USA INC

10MG

A211897 001 Aug 18, 2022

20MG

A211897 002 Aug 18, 2022

30MG

A211897 003 Aug 18, 2022

UNICHEM

10MG

A211819 001 Feb 17, 2021

20MG

A211819 002 Feb 17, 2021

30MG

A211819 003 Feb 17, 2021

APREPITANT

CAPSULE;ORAL

EMEND

+ MERCK

40MG **

N021549 003 Jun 30, 2006

ARBUTAMINE HYDROCHLORIDE

INJECTABLE;INJECTION

GENESA

GENSIA AUTOMEDICS

0.05MG/ML

N020420 001 Sep 12, 1997

ARDEPARIN SODIUM

INJECTABLE;INJECTION

NORMIFLO

+ PHARMACIA AND UPJOHN

5,000 UNITS/0.5ML **

N020227 002 May 23, 1997

+

10,000 UNITS/0.5ML **

N020227 001 May 23, 1997

ARFORMOTEROL TARTRATE

SOLUTION;INHALATION

ARFORMOTEROL TARTRATE

GLENMARK PHARMS LTD EQ 0.015MG BASE/2ML

A213132 001 Jun 22, 2021

ARGATROBAN

INJECTABLE;INJECTION

ARGATROBAN

+ SANDOZ

250MG/2.5ML (100MG/ML)

N020883 001 Jun 30, 2000

INJECTABLE;INTRAVENOUS

ARGATROBAN IN 0.9% SODIUM CHLORIDE

TEVA PHARMS USA

250MG/250ML (1MG/ML)

N206769 001 Dec 15, 2014

ARGATROBAN IN SODIUM CHLORIDE

+ CIPLA

50MG/50ML (1MG/ML) **

N022434 001 Jun 29, 2011

SOLUTION;INTRAVENOUS

ARGATROBAN IN DEXTROSE

SANDOZ

125MG/125ML (1MG/ML)

N201743 001 May 09, 2011

ARIPIPRAZOLE

INJECTABLE;INTRAMUSCULAR

ABILIFY

OTSUKA

9.75MG/1.3ML (7.5MG/ML)

N021866 001 Sep 20, 2006

SOLUTION;ORAL

ABILIFY

+ OTSUKA

1MG/ML **

N021713 001 Dec 10, 2004

TABLET;ORAL

ABILIFY MYCITE KIT

+ OTSUKA

2MG

N207202 001 Nov 13, 2017

+

5MG

N207202 002 Nov 13, 2017

+

10MG

N207202 003 Nov 13, 2017

+

15MG

N207202 004 Nov 13, 2017

+

20MG

N207202 005 Nov 13, 2017

+

30MG

N207202 006 Nov 13, 2017

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ARIPIPRAZOLE

TABLET; ORAL

ARIPIPRAZOLE

AIPING PHARM INC	2MG	A091279 001	Jan 09, 2017
	5MG	A091279 002	Jan 09, 2017
	10MG	A091279 003	Jan 09, 2017
	15MG	A091279 004	Jan 09, 2017
	20MG	A091279 005	Jan 09, 2017
	30MG	A091279 006	Jan 09, 2017
NORVIUM BIOSCIENCE	2MG	A206240 001	Sep 19, 2018
	5MG	A206240 002	Sep 19, 2018
	10MG	A206240 003	Sep 19, 2018
	15MG	A206240 004	Sep 19, 2018
	20MG	A206240 005	Sep 19, 2018
	30MG	A206240 006	Sep 19, 2018
TEVA PHARMS USA	2MG	A078607 001	Apr 28, 2015
	5MG	A078607 002	Apr 28, 2015
	10MG	A078608 001	Apr 28, 2015
	15MG	A078708 001	Apr 28, 2015
	20MG	A078708 002	Apr 28, 2015
	30MG	A078708 003	Apr 28, 2015
ZYDUS PHARMS	2MG	A090472 001	Jan 07, 2019
	5MG	A090472 002	Jan 07, 2019
	10MG	A090472 003	Jan 07, 2019
	15MG	A090472 004	Jan 07, 2019
	20MG	A090472 005	Jan 07, 2019
	30MG	A090472 006	Jan 07, 2019

TABLET, ORALLY DISINTEGRATING; ORAL

ABILIFY

+ OTSUKA	10MG **	N021729 002	Jun 07, 2006
+	15MG **	N021729 003	Jun 07, 2006
+	20MG **	N021729 004	Jun 07, 2006
+	30MG **	N021729 005	Jun 07, 2006

ARMODAFINIL

TABLET; ORAL

ARMODAFINIL

COREPHARMA	50MG	A201514 001	Mar 25, 2019
	150MG	A201514 002	Mar 25, 2019
	250MG	A201514 003	Mar 25, 2019
WATSON LABS INC	100MG	A200156 002	Aug 29, 2012
	200MG	A200156 004	Aug 29, 2012
NUVIGIL			
+ CEPHALON	100MG **	N021875 002	Mar 26, 2009

ARSENIC TRIOXIDE

INJECTABLE; INJECTION

ARSENIC TRIOXIDE

EUGIA PHARMA	2MG/ML	A214011 001	Oct 15, 2021
INGENUS PHARMS LLC	2MG/ML	A209315 002	Jan 14, 2021
TRISENOX			
+ CEPHALON	1MG/ML **	N021248 001	Sep 25, 2000

ARTICAINE HYDROCHLORIDE; EPINEPHRINE BITARTRATE

INJECTABLE; INJECTION

ARTICAINE HYDROCHLORIDE AND EPINEPHRINE BITARTRATE

HOSPIRA	4%;EQ 0.017MG BASE/1.7ML (4%;EQ 0.01MG BASE/ML)	A079138 001	Jun 18, 2010
ULTACAN			
HANSAMED INC	4%;EQ 0.0085MG BASE/1.7ML (4%; EQ 0.005MG BASE/ML)	A201751 001	Jul 11, 2017
ULTACAN FORTE			
HANSAMED INC	4%;EQ 0.017MG BASE/1.7ML (4%; EQ 0.01MG BASE/ML)	A201750 001	Jul 11, 2017

ASCORBIC ACID; BIOTIN; CYANOCOBALAMIN; DEXPANTHENOL; ERGOCALCIFEROL; FOLIC ACID; NIACINAMIDE; PHYTONADIONE; PYRIDOXINE HYDROCHLORIDE; RIBOFLAVIN 5'-PHOSPHATE SODIUM; THIAMINE HYDROCHLORIDE; VITAMIN A; VITAMIN E

FOR SOLUTION; INTRAVENOUS

M.V.I. PEDIATRIC

+ HOSPIRA	80MG/VIAL;0.02MG/VIAL;0.001MG/VIAL;5MG/VIAL;0.01MG/VIAL;0.14MG/VIAL;	N018920 001	Sep 21, 2000
-----------	--	-------------	--------------

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ASCORBIC ACID; BIOTIN; CYANOCOBALAMIN; DEXPANTHENOL; ERGOCALCIFEROL; FOLIC ACID; NIACINAMIDE; PHYTONADIONE; PYRIDOXINE HYDROCHLORIDE; RIBOFLAVIN 5'-PHOSPHATE SODIUM; THIAMINE HYDROCHLORIDE; VITAMIN A; VITAMIN E

FOR SOLUTION; INTRAVENOUS
M.V.I. PEDIATRIC

17MG/VIAL; 0.2MG/VIAL; 1MG/VIAL; 1.4MG/VIAL;
EQ 1.2MG
BASE/VIAL; 0.7MG/VIAL; 7MG/VIAL

ASCORBIC ACID; BIOTIN; CYANOCOBALAMIN; DEXPANTHENOL; ERGOCALCIFEROL; FOLIC ACID; NIACINAMIDE; PYRIDOXINE HYDROCHLORIDE; RIBOFLAVIN 5'-PHOSPHATE SODIUM; THIAMINE HYDROCHLORIDE; VITAMIN A PALMITATE; VITAMIN E

INJECTABLE; INJECTION
BEROCCA PN

ROCHE 50MG/ML; 0.03MG/ML; 0.0025MG/ML; 7.5MG/ML; N006071 003 Oct 10, 1985
100
IU/ML; 0.2MG/ML; 20MG/ML; 2MG/ML; 1.8MG/ML;

ASCORBIC ACID; BIOTIN; CYANOCOBALAMIN; DEXPANTHENOL; ERGOCALCIFEROL; FOLIC ACID; NIACINAMIDE; PYRIDOXINE HYDROCHLORIDE; RIBOFLAVIN 5'-PHOSPHATE SODIUM; THIAMINE HYDROCHLORIDE; VITAMIN A; VITAMIN E

INJECTABLE; INJECTION
M.V.C. 9+3

ABRAXIS PHARM 10MG/ML; 0.006MG/ML; 0.5MCG/ML; 1.5MG/ML; 2 N018440 002 Aug 08, 1985
0
IU/ML; 0.04MG/ML; 4MG/ML; 0.4MG/ML; 0.36MG/

M.V.I.-12 ADULT
HOSPIRA

10MG/ML; 0.006MG/ML; 0.5MCG/ML; 1.5MG/ML; 2 N008809 004 Aug 08, 1985
0
IU/ML; 0.04MG/ML; 4MG/ML; 0.4MG/ML; 0.36MG/

+

20MG/ML; 0.006MG/ML; 0.05MCG/ML; 1.5MG/ML; N008809 006 Sep 09, 2004
0.0005MG/ML; 0.06MG/ML; 4MG/ML; 0.6MG/ML; 0.36MG/ML; 0.6MG/ML; 0.1MG/ML; 1MG/ML

MVC PLUS

WATSON LABS 10MG/ML; 0.006MG/ML; 0.5MCG/ML; 1.5MG/ML; 2 N018439 002 Aug 08, 1985
0
IU/ML; 0.04MG/ML; 4MG/ML; 0.4MG/ML; 0.36MG/

ASCORBIC ACID; BIOTIN; CYANOCOBALAMIN; DEXPANTHENOL; ERGOCALCIFEROL; FOLIC ACID; NIACINAMIDE; PYRIDOXINE HYDROCHLORIDE; RIBOFLAVIN 5'-PHOSPHATE SODIUM; THIAMINE HYDROCHLORIDE; VITAMIN A; VITAMIN E; VITAMIN K

INJECTABLE; INTRAVENOUS
M.V.I. ADULT

+ HOSPIRA

200MG/VIAL; 0.06MG/VIAL; 0.005MG/VIAL; 15M N021625 001 Jan 30, 2004
G/VIAL; 0.005MG/VIAL; 0.6MG/VIAL; 40MG/VIAL;
6MG/VIAL; 3.6MG/VIAL; 6MG/VIAL; 1MG/VIAL;
10MG/VIAL; 0.15MG/VIAL

M.V.I. ADULT (PHARMACY BULK PACKAGE)

+ HOSPIRA

200MG/5ML; 0.06MG/5ML; 0.005MG/5ML; 15MG/5 N021643 001 Feb 18, 2004
ML; 0.005MG/5ML; 0.6MG/5ML; 40MG/5ML; 6MG/5
ML; 3.6MG/5ML; 6MG/5ML; 1MG/5ML; 10MG/5ML; 0.15MG/5ML

ASCORBIC ACID; BIOTIN; CYANOCOBALAMIN; DEXPANTHENOL; ERGOCALCIFEROL; FOLIC ACID; NIACINAMIDE; PYRIDOXINE HYDROCHLORIDE; RIBOFLAVIN; THIAMINE HYDROCHLORIDE; VITAMIN A; VITAMIN E

INJECTABLE; INJECTION
M.V.I.-12 ADULT

HOSPIRA

20MG/ML; 0.006MG/ML; 0.5MCG/ML; 1.5MG/ML; 2 N008809 005 Apr 22, 2004
0
IU/ML; 0.6MG/ML; 4MG/ML; 0.4MG/ML; 0.36MG/M

ASCORBIC ACID; BIOTIN; CYANOCOBALAMIN; DEXPANTHENOL; ERGOCALCIFEROL; FOLIC ACID; NIACINAMIDE; PYRIDOXINE; RIBOFLAVIN 5'-PHOSPHATE SODIUM; THIAMINE; VITAMIN A; VITAMIN E

INJECTABLE; INJECTION
M.V.I.-12 LYOPHILIZED

TELIGENT

100MG/VIAL; 0.06MG/VIAL; 0.005MG/VIAL; 15M N018933 002 Aug 08, 1985
G/VIAL; 5MCG/VIAL; 0.4MG/VIAL; 40MG/VIAL; 4
MG/VIAL; 3.6MG/VIAL; 3MG/VIAL; 1MG/VIAL; 10
MG/VIAL

ASCORBIC ACID; BIOTIN; CYANOCOBALAMIN; ERGOCALCIFEROL; FOLIC ACID; NIACINAMIDE; PANTOTHENIC ACID; PHYTONADIONE; PYRIDOXINE; RIBOFLAVIN; THIAMINE; VITAMIN A PALMITATE; VITAMIN E

INJECTABLE; INJECTION
VITAPED

HOSPIRA

N/A, 80MG/VIAL; N/A, 0.02MG/VIAL; N/A, 0.001 N020176 001 Dec 29, 1993
MG/VIAL; 400
IU/10ML, N/A, N/A, 0.14MG/VIAL; N/A, 17MG/VIAL;
N/A, 5MG/VIAL; 0.2MG/10ML, N/A, N/A, 1MG/

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ASCORBIC ACID; BIOTIN; CYANOCOBALAMIN; ERGOCALCIFEROL; FOLIC ACID; NIACINAMIDE; PANTOTHENIC ACID; PHYTONADIONE; PYRIDOXINE; RIBOFLAVIN; THIAMINE; VITAMIN A PALMITATE; VITAMIN E INJECTABLE; INJECTION VITAPED

MG/VIAL;N/A,1.2MG/VIAL;EQ 2,300 UNITS
BASE/10ML,N/A;7 IU/10ML,N/A

ASPIRIN

CAPSULE, EXTENDED RELEASE;ORAL

DURLAZA

+ HESP 162.5MG N200671 001 Sep 04, 2015

TABLET;ORAL

BAYER EXTRA STRENGTH ASPIRIN FOR MIGRAINE PAIN

BAYER 500MG N021317 001 Oct 18, 2001

TABLET, EXTENDED RELEASE;ORAL

8-HOUR BAYER

BAYER 650MG N016030 001

MEASURIN

BAYER 650MG N016030 002

ASPIRIN; BUTALBITAL

TABLET;ORAL

AXOTAL

SAVAGE LABS 650MG;50MG A088305 001 Oct 13, 1983

ASPIRIN; BUTALBITAL; CAFFEINE

CAPSULE;ORAL

BUTALBITAL, ASPIRIN AND CAFFEINE

LGM PHARMA 500MG;50MG;40MG A205230 001 Oct 18, 2021

WATSON LABS 325MG;50MG;40MG A086231 002 Feb 12, 1985

FIORINAL

+ ALLERGAN 325MG;50MG;40MG ** N017534 005 Apr 16, 1986

TABLET;ORAL

BUTALBITAL, ASPIRIN AND CAFFEINE

ACTAVIS ELIZABETH 325MG;50MG;40MG A086710 002 Aug 23, 1983

FOSUN PHARMA 325MG;50MG;40MG A086398 002 Apr 06, 1984

HALSEY 325MG;50MG;40MG A089448 001 Dec 01, 1986

+ HIKMA INTL PHARMS 325MG;50MG;40MG A086162 002 Feb 16, 1984

IVAX PHARMS 325MG;50MG;40MG A085441 002 Oct 31, 1984

PURACAP PHARM 325MG;50MG;40MG A087048 002 Dec 09, 1983

QUANTUM PHARMICS 325MG;50MG;40MG A088972 001 Jun 18, 1985

WATSON LABS 325MG;50MG;40MG A086237 002 Mar 23, 1984

FIORINAL

+ ALLERGAN 325MG;50MG;40MG ** N017534 003 Apr 16, 1986

LANORINAL

LANNETT 325MG;50MG;40MG A086986 002 Oct 18, 1985

ASPIRIN; BUTALBITAL; CAFFEINE; CODEINE PHOSPHATE

CAPSULE;ORAL

BUTALBITAL, ASPIRIN, CAFFEINE, AND CODEINE PHOSPHATE

DR REDDYS LABS SA 325MG;50MG;40MG;30MG A203335 001 Oct 30, 2015

ESJAY PHARMA 325MG;50MG;40MG;30MG A075351 001 Mar 05, 1999

WATSON LABS 325MG;50MG;40MG;30MG A074359 001 Aug 31, 1995

FIORINAL W/CODEINE

+ ALLERGAN 325MG;50MG;40MG;30MG N019429 003 Oct 26, 1990

ASPIRIN; CAFFEINE; DIHYDROCODEINE BITARTRATE

CAPSULE;ORAL

SYNALGOS-DC

+ SUN PHARM INDUSTRIES 356.4MG;30MG;16MG N011483 004 Sep 06, 1983

ASPIRIN; CAFFEINE; ORPHENADRINE CITRATE

TABLET;ORAL

INVAGESIC

CHARTWELL RX 385MG;30MG;25MG A074817 001 Nov 27, 1996

INVAGESIC FORTE

CHARTWELL RX 770MG;60MG;50MG A074817 002 Nov 27, 1996

NORGESIC

+ BAUSCH 385MG;30MG;25MG ** N013416 003 Oct 27, 1982

NORGESIC FORTE

+ BAUSCH 770MG;60MG;50MG ** N013416 004 Oct 27, 1982

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ASPIRIN; CAFFEINE; ORPHENADRINE CITRATE

TABLET; ORAL

ORPHENADRINE CITRATE, ASPIRIN, AND CAFFEINE

SANDOZ	385MG; 30MG; 25MG	A074654 001	Dec 31, 1996
	770MG; 60MG; 50MG	A074654 002	Dec 31, 1996
STEVENS J	385MG; 30MG; 25MG	A074988 001	Apr 30, 1999
	770MG; 60MG; 50MG	A074988 002	Apr 30, 1999

ASPIRIN; CAFFEINE; PROPOXYPHENE HYDROCHLORIDE

CAPSULE; ORAL

COMPOUND 65

ALRA	389MG; 32.4MG; 65MG	A084553 002	Aug 17, 1983
DARVON COMPOUND			
XANODYNE PHARM	389MG; 32.4MG; 32MG	N010996 006	Mar 08, 1983
DARVON COMPOUND-65			
XANODYNE PHARM	389MG; 32.4MG; 65MG	N010996 007	Mar 08, 1983
PROPOXYPHENE COMPOUND 65			
IVAX SUB TEVA PHARMS	389MG; 32.4MG; 65MG	A083077 002	Dec 07, 1984
+ SANDOZ	389MG; 32.4MG; 65MG	A080044 002	Sep 16, 1983
TEVA	389MG; 32.4MG; 65MG	A089025 001	Mar 29, 1985
PROPOXYPHENE COMPOUND-65			
SANDOZ	389MG; 32.4MG; 65MG	A083101 002	Jun 24, 1985
PROPOXYPHENE HYDROCHLORIDE W/ ASPIRIN AND CAFFEINE			
WATSON LABS	389MG; 32.4MG; 65MG	A085732 002	Sep 03, 1984

ASPIRIN; CARISOPRODOL

TABLET; ORAL

CARISOPRODOL AND ASPIRIN

CHARTWELL RX	325MG; 200MG	A089594 001	Mar 31, 1989
GENUS	325MG; 200MG	A040116 001	Apr 25, 1996
NOVAST LABS	325MG; 200MG	A040832 001	Jan 07, 2010
OXFORD PHARMS	325MG; 200MG	A040252 001	Dec 10, 1997
CARISOPRODOL COMPOUND			
WATSON LABS	325MG; 200MG	A088809 001	Oct 03, 1985
SOMA COMPOUND			
MEDA PHARMS	325MG; 200MG **	N012365 005	Jul 11, 1983

ASPIRIN; CARISOPRODOL; CODEINE PHOSPHATE

TABLET; ORAL

CARISOPRODOL, ASPIRIN AND CODEINE PHOSPHATE

INGENUS PHARMS NJ	325MG; 200MG; 16MG	A040860 001	Jan 07, 2010
OXFORD PHARMS	325MG; 200MG; 16MG	A040283 001	Dec 29, 1998
SANDOZ	325MG; 200MG; 16MG	A040118 001	Apr 16, 1996
SOMA COMPOUND W/ CODEINE			
MEDA PHARMS	325MG; 200MG; 16MG **	N012366 002	Jul 11, 1983

ASPIRIN; DIPYRIDAMOLE

CAPSULE, EXTENDED RELEASE; ORAL

AGGRENOL

+ BOEHRINGER INGELHEIM	25MG; 200MG **	N020884 001	Nov 22, 1999
ASPIRIN AND DIPYRIDAMOLE			
ANI PHARMS	25MG; 200MG	A206964 001	Jan 18, 2017
CHARTWELL MOLECULAR	25MG; 200MG	A204552 001	Mar 20, 2019
SUN PHARM	25MG; 200MG	A208572 001	Aug 21, 2018

ASPIRIN; HYDROCODONE BITARTRATE

TABLET; ORAL

AZDONE

SCHWARZ PHARMA	500MG; 5MG **	A089420 001	Jan 25, 1988
VICOPRIN			
ABBOTT	500MG; 5MG	A086333 001	Sep 14, 1983

ASPIRIN; MEPROBAMATE

TABLET; ORAL

EQUAGESIC

SUN PHARM INDUSTRIES	325MG; 200MG	N011702 003	Dec 29, 1983
MEPRO-ASPIRIN			
SANDOZ	325MG; 200MG	A089127 001	Mar 02, 1987
MEPROBAMATE AND ASPIRIN			
PAR PHARM	325MG; 200MG	A089126 001	Aug 19, 1986
MICRAININ			
MEDPOINTE PHARM HLC	325MG; 200MG	A084978 001	

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ASPIRIN; MEPROBAMATE

TABLET; ORAL

Q-GESIC

QUANTUM PHARMICS 325MG;200MG

A088740 001 Jun 01, 1984

ASPIRIN; METHOCARBAMOL

TABLET; ORAL

METHOCARBAMOL AND ASPIRIN

IVAX SUB TEVA PHARMS 325MG;400MG

A087211 001 Dec 22, 1982

MCNEIL 325MG;400MG

A089193 001 Feb 12, 1986

PAR PHARM 325MG;400MG

A089657 001 Nov 04, 1988

STEVENS J 325MG;400MG

A081145 001 Jan 31, 1995

ROBAXISAL

+ ROBINS AH 325MG;400MG

N012281 001

ASPIRIN; OMEPRAZOLE

TABLET, DELAYED RELEASE; ORAL

YOSPRALA

+ GENUS LIFESCIENCES 81MG;40MG

N205103 001 Sep 14, 2016

+ 325MG;40MG

N205103 002 Sep 14, 2016

ASPIRIN; OXYCODONE HYDROCHLORIDE

TABLET; ORAL

OXYCODONE AND ASPIRIN

ACTAVIS LABS FL INC 325MG;4.8355MG

A090084 001 Mar 22, 2011

DR REDDYS LABS SA 325MG;4.8355MG

A091670 001 Mar 16, 2011

ASPIRIN; OXYCODONE HYDROCHLORIDE; OXYCODONE TEREPHTHALATE

TABLET; ORAL

CODOXY

HALSEY 325MG;4.5MG;0.38MG

A087464 001 Jul 01, 1982

OXYCODONE AND ASPIRIN

ANI PHARMS 325MG;4.5MG;0.38MG

A040255 001 Feb 27, 1998

SUN PHARM INDUSTRIES 325MG;4.5MG;0.38MG

A040260 001 Jul 17, 1998

325MG;4.5MG;0.38MG

A087794 001 May 26, 1982

OXYCODONE AND ASPIRIN (HALF-STRENGTH)

ROXANE 325MG;2.25MG;0.19MG

A087742 001 Jun 04, 1982

PERCODAN

ENDO OPERATIONS 325MG;4.5MG;0.38MG **

N007337 006

PERCODAN-DEMI

ENDO OPERATIONS 325MG;2.25MG;0.19MG **

N007337 005

ROXIPRIN

ROXANE 325MG;4.5MG;0.38MG

A087743 001 Jun 04, 1982

ASPIRIN; PENTAZOCINE HYDROCHLORIDE

TABLET; ORAL

TALWIN COMPOUND

+ SANOFI AVENTIS US 325MG;EQ 12.5MG BASE **

N016891 001

ASPIRIN; PRAVASTATIN SODIUM

TABLET; ORAL

PRAVIGARD PAC (COPACKAGED)

BRISTOL MYERS SQUIBB 81MG;20MG

N021387 001 Jun 24, 2003

81MG;40MG

N021387 002 Jun 24, 2003

81MG;80MG

N021387 003 Jun 24, 2003

325MG;20MG

N021387 004 Jun 24, 2003

325MG;40MG

N021387 005 Jun 24, 2003

325MG;80MG

N021387 006 Jun 24, 2003

ASPIRIN; PROPOXYPHENE HYDROCHLORIDE

CAPSULE; ORAL

DARVON W/ ASA

XANODYNE PHARM 325MG;65MG

N010996 005

ASPIRIN; PROPOXYPHENE NAPSYLATE

CAPSULE; ORAL

DARVON-N W/ ASA

AAIPHARMA LLC 325MG;100MG

N016829 001

TABLET; ORAL

DARVON-N W/ ASA

AAIPHARMA LLC 325MG;100MG

N016863 001

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ATAZANAVIR SULFATE

CAPSULE;ORAL

ATAZANAVIR SULFATE

AMNEAL	EQ 100MG BASE	A209717 001	Jun 01, 2020
	EQ 150MG BASE	A209717 002	Jun 01, 2020
	EQ 200MG BASE	A209717 003	Jun 01, 2020
	EQ 300MG BASE	A209717 004	Jun 01, 2020
CIPLA	EQ 100MG BASE	A200626 001	Aug 09, 2018
	EQ 150MG BASE	A200626 002	Aug 09, 2018
	EQ 200MG BASE	A200626 003	Aug 09, 2018
	EQ 300MG BASE	A200626 004	Aug 09, 2018
MYLAN	EQ 150MG BASE	A208177 001	Sep 24, 2018
	EQ 200MG BASE	A208177 002	Sep 24, 2018
	EQ 300MG BASE	A208177 003	Sep 24, 2018
ZYDUS PHARMS	EQ 150MG BASE	A210575 001	Jun 04, 2020
	EQ 200MG BASE	A210575 002	Jun 04, 2020
	EQ 300MG BASE	A210575 003	Jun 04, 2020
REYATAZ			
+	BRISTOL MYERS SQUIBB	EQ 100MG BASE **	N021567 001 Jun 20, 2003
+		EQ 150MG BASE **	N021567 002 Jun 20, 2003

ATENOLOL

INJECTABLE;INJECTION

TENORMIN

+ ASTRAZENECA

0.5MG/ML **

N019058 001 Sep 13, 1989

TABLET;ORAL

ATENOLOL

ABLE	25MG	A076907 001	Jul 30, 2004
	50MG	A076907 002	Jul 30, 2004
	100MG	A076907 003	Jul 30, 2004
APOTHECON	50MG	A073317 001	Mar 20, 1992
	100MG	A073318 001	Mar 20, 1992
CHARTWELL RX	25MG	A074265 001	Feb 28, 1994
	50MG	A074265 002	Feb 28, 1994
	100MG	A074265 003	Feb 28, 1994
ESJAY PHARMA	25MG	A074099 001	Apr 28, 1992
	50MG	A073542 001	Dec 19, 1991
	100MG	A073543 001	Dec 19, 1991
IPCA LABS LTD	25MG	A077877 001	Dec 27, 2006
	50MG	A077877 002	Dec 27, 2006
	100MG	A077877 003	Dec 27, 2006
MYLAN	25MG	A074126 003	Aug 26, 1998
	50MG	A074126 001	Mar 23, 1994
	100MG	A074126 002	Mar 23, 1994
NORTHSTAR HLTHCARE	25MG	A078254 001	Sep 25, 2009
	50MG	A078254 002	Sep 25, 2009
	100MG	A078254 003	Sep 25, 2009
NOSTRUM LABS	50MG	A074127 001	Feb 21, 1995
	100MG	A074127 002	Feb 21, 1995
PLIVA	25MG	A074101 001	Jul 17, 1997
	50MG	A074101 002	Jul 17, 1997
	100MG	A074101 003	Jul 17, 1997
SCS	50MG	A073676 001	Oct 30, 1992
	100MG	A073676 002	Oct 30, 1992
SUN PHARM INDS INC	25MG	A078210 001	Jul 10, 2007
	50MG	A078210 002	Jul 10, 2007
	100MG	A078210 003	Jul 10, 2007
SUN PHARM INDUSTRIES	25MG	A074499 001	Jul 30, 1997
	50MG	A073475 001	Mar 30, 1993
	100MG	A073476 001	Mar 30, 1993
TEVA	50MG	A073315 001	May 28, 1993
	100MG	A073316 001	May 28, 1993
TEVA PHARMS	50MG	A074120 001	Feb 24, 1995
	100MG	A074120 002	Feb 24, 1995
WATSON LABS	50MG	A073352 001	Dec 27, 1991
WATSON LABS TEVA	100MG	A073353 001	Dec 27, 1991

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ATENOLOL; CHLORTHALIDONE

TABLET; ORAL

ATENOLOL AND CHLORTHALIDONE

AUROBINDO PHARMA USA	50MG;25MG	A074203	001	Oct 31, 1993
	100MG;25MG	A074203	002	Oct 31, 1993
NOSTRUM LABS	50MG;25MG	A074404	001	May 14, 1998
	100MG;25MG	A074404	002	May 14, 1998
PLIVA	50MG;25MG	A074107	001	Sep 24, 1997
	100MG;25MG	A074107	002	Sep 24, 1997
SUN PHARM INDUSTRIES	50MG;25MG	A073582	002	Apr 29, 1993
	100MG;25MG	A073582	001	Apr 29, 1993

ATOMOXETINE HYDROCHLORIDE

CAPSULE; ORAL

STRATTERA

LILLY	EQ 5MG BASE	N021411	001	Nov 26, 2002
-------	-------------	---------	-----	--------------

ATORVASTATIN CALCIUM

TABLET; ORAL

ATORVASTATIN CALCIUM

CADILA PHARMS LTD	EQ 10MG BASE	A212103	001	Oct 16, 2023
	EQ 20MG BASE	A212103	002	Oct 16, 2023
	EQ 40MG BASE	A212103	003	Oct 16, 2023
	EQ 80MG BASE	A212103	004	Oct 16, 2023
INVAGEN PHARMS	EQ 10MG BASE	A204846	001	Jan 09, 2017
	EQ 20MG BASE	A204846	002	Jan 09, 2017
	EQ 40MG BASE	A204846	003	Jan 09, 2017
	EQ 80MG BASE	A204846	004	Jan 09, 2017
LEPU PHARM	EQ 10MG BASE	A216848	001	Nov 03, 2022
	EQ 20MG BASE	A216848	002	Nov 03, 2022
	EQ 40MG BASE	A216848	003	Nov 03, 2022
	EQ 80MG BASE	A216848	004	Nov 03, 2022
LUPIN LTD	EQ 10MG BASE	A204991	001	Mar 06, 2019
	EQ 20MG BASE	A204991	002	Mar 06, 2019
	EQ 40MG BASE	A204991	003	Mar 06, 2019
	EQ 80MG BASE	A204991	004	Mar 06, 2019
PERRIGO R AND D	EQ 10MG BASE	A208478	001	Jun 23, 2020
	EQ 20MG BASE	A208478	002	Jun 23, 2020
STRIDES PHARMA	EQ 10MG BASE	A217634	001	Dec 21, 2023
	EQ 20MG BASE	A217634	002	Dec 21, 2023
	EQ 40MG BASE	A217634	003	Dec 21, 2023
	EQ 80MG BASE	A217634	004	Dec 21, 2023
TEVA PHARMS	EQ 10MG BASE	A078773	001	May 29, 2012
	EQ 20MG BASE	A078773	002	May 29, 2012
	EQ 40MG BASE	A078773	003	May 29, 2012
	EQ 80MG BASE	A078773	004	May 29, 2012

ATORVASTATIN CALCIUM; EZETIMIBE

TABLET; ORAL

LIPTRUZET

+	ORGANON	EQ 10MG BASE;10MG **	N200153	001	May 03, 2013
+		EQ 20MG BASE;10MG **	N200153	002	May 03, 2013
+		EQ 40MG BASE;10MG **	N200153	003	May 03, 2013
+		EQ 80MG BASE;10MG **	N200153	004	May 03, 2013

ATOVAQUONE

TABLET; ORAL

MEPRON

+	GLAXOSMITHKLINE LLC	250MG **	N020259	001	Nov 25, 1992
---	---------------------	----------	---------	-----	--------------

ATRACURIUM BESYLATE

INJECTABLE; INJECTION

ATRACURIUM BESYLATE

BAXTER HLTHCARE	10MG/ML	A074824	001	Sep 30, 1997
BAXTER HLTHCARE CORP	10MG/ML	A074753	001	Jan 23, 1997
HOSPIRA	10MG/ML	A074632	001	Dec 23, 1996
	10MG/ML	A074740	001	Mar 28, 1997
NORVIUM BIOSCIENCE	10MG/ML	A206096	001	Jun 22, 2017
TEVA PARENTERAL	10MG/ML	A074784	001	Jun 11, 1997
WATSON PHARMS TEVA	10MG/ML	A074945	001	Jul 28, 1998

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ATRACURIUM BESYLATE

INJECTABLE; INJECTION

ATRACURIUM BESYLATE PRESERVATIVE FREE

BAXTER HLTHCARE	10MG/ML	A074825	001	Sep 30, 1997
BAXTER HLTHCARE CORP	10MG/ML	A074768	001	Jan 23, 1997
HOSPIRA	10MG/ML	A074633	001	Dec 23, 1996
	10MG/ML	A074639	001	Mar 25, 1997
	10MG/ML	A074741	001	Mar 28, 1997
NORVIUM BIOSCIENCE	10MG/ML	A206001	001	Apr 07, 2017
WATSON LABS INC	10MG/ML	A074944	001	Jul 28, 1998

TRACRIUM

+ HOSPIRA	10MG/ML **	N018831	002	Jun 20, 1985
TRACRIUM PRESERVATIVE FREE				
+ HOSPIRA	10MG/ML **	N018831	001	Nov 23, 1983

ATROPINE

SOLUTION; INTRAMUSCULAR

ATROPEN

+ MMT	EQ 0.25MG SULFATE/0.3ML	N017106	004	Sep 17, 2004
+	EQ 0.5MG SULFATE/0.7ML	N017106	003	Jun 19, 2003
+	EQ 1MG SULFATE/0.7ML	N017106	002	Jun 19, 2003
+	EQ 2MG SULFATE/0.7ML	N017106	001	

ATROPINE

ABBVIE	EQ 2MG SULFATE/0.7ML	A071295	001	Jan 30, 1987
ATROPINE (AUTOINJECTOR)				
RAFA LABS LTD	EQ 2MG SULFATE/0.7ML (EQ 2MG SULFATE/0.7ML)	N212319	001	Jul 09, 2018

ATROPINE SULFATE

AEROSOL, METERED; INHALATION

ATROPINE SULFATE

US ARMY	EQ 0.36MG BASE/INH	N020056	001	Sep 19, 1990
---------	--------------------	---------	-----	--------------

ATROPINE SULFATE; DIFENOXIN HYDROCHLORIDE

TABLET; ORAL

MOTOFEN HALF-STRENGTH

LEGACY PHARMA	0.025MG; 0.5MG	N017744	001	
---------------	----------------	---------	-----	--

ATROPINE SULFATE; DIPHENOXYLATE HYDROCHLORIDE

CAPSULE; ORAL

DIPHENOXYLATE HYDROCHLORIDE W/ ATROPINE SULFATE

SCHERER RP	0.025MG; 2.5MG	A086440	001	
------------	----------------	---------	-----	--

SOLUTION; ORAL

COLONAIID

MEDPOINTE PHARM HLC	0.025MG/5ML; 2.5MG/5ML	A085735	001	
---------------------	------------------------	---------	-----	--

LOMANATE

ALPHARMA US PHARMS	0.025MG/5ML; 2.5MG/5ML	A085746	001	
--------------------	------------------------	---------	-----	--

LOMOTIL

GD SEARLE LLC	0.025MG/5ML; 2.5MG/5ML	N012699	001	
---------------	------------------------	---------	-----	--

TABLET; ORAL

COLONAIID

MEDPOINTE PHARM HLC	0.025MG; 2.5MG	A085737	001	
---------------------	----------------	---------	-----	--

DI-ATRO

MD PHARM	0.025MG; 2.5MG	A085266	001	
----------	----------------	---------	-----	--

DIPHENOXYLATE HYDROCHLORIDE AND ATROPINE SULFATE

ABLE	0.025MG; 2.5MG	A040395	001	Nov 27, 2000
------	----------------	---------	-----	--------------

ANI PHARMS	0.025MG; 2.5MG	A086727	001	
------------	----------------	---------	-----	--

ASCOT	0.025MG; 2.5MG	A087934	001	Jul 19, 1983
-------	----------------	---------	-----	--------------

DR REDDYS LABS SA	0.025MG; 2.5MG	A210789	001	Jun 03, 2020
-------------------	----------------	---------	-----	--------------

FOSUN PHARMA	0.025MG; 2.5MG	A086173	001	
--------------	----------------	---------	-----	--

HEATHER	0.025MG; 2.5MG	A086798	001	
---------	----------------	---------	-----	--

HIKMA	0.025MG; 2.5MG	A087765	001	Mar 15, 1982
-------	----------------	---------	-----	--------------

INWOOD LABS	0.025MG; 2.5MG	A085509	001	
-------------	----------------	---------	-----	--

KV PHARM	0.025MG; 2.5MG	A085659	001	
----------	----------------	---------	-----	--

LEDERLE	0.025MG; 2.5MG	A086950	001	
---------	----------------	---------	-----	--

MYLAN	0.025MG; 2.5MG	A085762	001	
-------	----------------	---------	-----	--

PARKE DAVIS	0.025MG; 2.5MG	A087131	001	
-------------	----------------	---------	-----	--

PVT FORM	0.025MG; 2.5MG	A085766	001	
----------	----------------	---------	-----	--

R AND S PHARMA	0.025MG; 2.5MG	A085035	001	
----------------	----------------	---------	-----	--

ROXANE	0.025MG; 2.5MG	A086057	001	
--------	----------------	---------	-----	--

STRIDES PHARMA	0.025MG; 2.5MG	A040357	001	May 02, 2000
----------------	----------------	---------	-----	--------------

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ATROPINE SULFATE; DIPHENOXYLATE HYDROCHLORIDE

TABLET; ORAL

DIPHENOXYLATE HYDROCHLORIDE AND ATROPINE SULFATE

SUN PHARM INDUSTRIES	0.025MG;2.5MG	A085506	001	
UPSHER SMITH LABS	0.025MG;2.5MG	A210571	001	Aug 31, 2018
USL PHARMA	0.025MG;2.5MG	A087842	001	Mar 29, 1982
VALEANT PHARM INTL	0.025MG;2.5MG	A087195	001	Feb 16, 1982
WATSON LABS	0.025MG;2.5MG	A085876	001	
LO-TROL				
VANGARD	0.025MG;2.5MG	A088009	001	Mar 25, 1983
LOGEN				
SUPERPHARM	0.025MG;2.5MG	A088962	001	May 10, 1985
LONOX				
FOSUN PHARMA	0.025MG;2.5MG	A085311	002	
LOW-QUEL				
HALSEY	0.025MG;2.5MG	A085211	001	

ATROPINE SULFATE; EDROPHONIUM CHLORIDE

INJECTABLE; INJECTION

ENLON-PLUS

NORVIUM BIOSCIENCE	0.14MG/ML;10MG/ML	N019677	001	Nov 06, 1991
+	0.14MG/ML;10MG/ML	N019678	001	Nov 06, 1991

ATROPINE SULFATE; MEPERIDINE HYDROCHLORIDE

INJECTABLE; INJECTION

ATROPINE AND DEMEROL

ABBVIE	0.4MG/ML;50MG/ML	A087853	001	Nov 26, 1982
	0.4MG/ML;75MG/ML	A087847	001	Nov 26, 1982
	0.4MG/ML;100MG/ML	A087848	001	Nov 26, 1982
MEPERIDINE AND ATROPINE SULFATE				
WYETH AYERST	0.4MG/ML;50MG/ML	A085121	001	
	0.4MG/ML;75MG/ML	A085121	002	
	0.4MG/ML;100MG/ML	A085121	003	

ATROPINE; PRALIDOXIME CHLORIDE

INJECTABLE; INTRAMUSCULAR

ATNAA

US ARMY	2.1MG/0.7ML;600MG/2ML	N021175	001	Jan 17, 2002
---------	-----------------------	---------	-----	--------------

AVOBENZONE; OCTINOXATE; OXYBENZONE

LOTION; TOPICAL

SHADE UVAGUARD

+	BAYER HEALTHCARE LLC	3%;7.5%;3%	N020045	001	Dec 07, 1992
---	----------------------	------------	---------	-----	--------------

AZACITIDINE

POWDER; INTRAVENOUS, SUBCUTANEOUS

AZACITIDINE

LUPIN LTD	100MG/VIAL	A210748	001	Feb 27, 2019
NORVIUM BIOSCIENCE	100MG/VIAL	A204949	001	Apr 28, 2016

AZATADINE MALEATE

TABLET; ORAL

OPTIMINE

SCHERING	1MG	N017601	001	
----------	-----	---------	-----	--

AZATADINE MALEATE; PSEUDOEPHEDRINE SULFATE

TABLET, EXTENDED RELEASE; ORAL

TRINALIN

SCHERING	1MG;120MG	N018506	001	Mar 23, 1982
----------	-----------	---------	-----	--------------

AZATHIOPRINE

TABLET; ORAL

IMURAN

+	LEGACY PHARMA	25MG **	N016324	002
---	---------------	---------	---------	-----

AZATHIOPRINE SODIUM

INJECTABLE; INJECTION

IMURAN

+	CASPER PHARMA LLC	EQ 100MG BASE/VIAL **	N017391	001
---	-------------------	-----------------------	---------	-----

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

AZITHROMYCIN

TABLET; ORAL

AZITHROMYCIN

ZYDUS PHARMS

EQ 250MG BASE

A213275 001 Apr 11, 2023

EQ 500MG BASE

A213056 001 Apr 07, 2023

ZITHROMAX

+ PFIZER

EQ 600MG BASE **

N050730 001 Jun 12, 1996

AZITHROMYCIN DIHYDRATE; TROVAFLOXACIN MESYLATE

FOR SUSPENSION, TABLET; ORAL

TROVAN/ZITHROMAX COMPLIANCE PAK

PFIZER

EQ 1GM BASE, N/A; N/A, EQ 100MG BASE

N050762 001 Dec 18, 1998

AZLOCILLIN SODIUM

INJECTABLE; INJECTION

AZLIN

BAYER PHARMS

EQ 2GM BASE/VIAL

A062388 001 Sep 08, 1982

EQ 2GM BASE/VIAL

A062417 001 Oct 12, 1982

EQ 2GM BASE/VIAL

N050562 001 Sep 03, 1982

EQ 3GM BASE/VIAL

A062388 002 Sep 08, 1982

EQ 3GM BASE/VIAL

A062417 002 Oct 12, 1982

EQ 3GM BASE/VIAL

N050562 002 Sep 03, 1982

EQ 4GM BASE/VIAL

A062388 003 Sep 08, 1982

EQ 4GM BASE/VIAL

A062417 003 Oct 12, 1982

EQ 4GM BASE/VIAL

N050562 003 Sep 03, 1982

AZTREONAM

INJECTABLE; INJECTION

AZACTAM

BRISTOL MYERS SQUIBB 500MG/VIAL **

N050580 001 Dec 31, 1986

AZACTAM IN PLASTIC CONTAINER

BRISTOL MYERS SQUIBB

10MG/ML

N050632 003 May 24, 1989

+

20MG/ML

N050632 002 May 24, 1989

+

40MG/ML

N050632 001 May 24, 1989

AZTREONAM

HIKMA

1GM/VIAL

A065286 001 Mar 23, 2011

2GM/VIAL

A065286 002 Mar 23, 2011

BACAMPICILLIN HYDROCHLORIDE

FOR SUSPENSION; ORAL

SPECTROBID

PFIZER

125MG/5ML

N050556 001 Mar 23, 1982

TABLET; ORAL

SPECTROBID

PFIZER

400MG

N050520 001

800MG

N050520 002 Sep 12, 1983

BACITRACIN

OINTMENT; OPHTHALMIC

BACIGUENT

PHARMACIA AND UPJOHN 500 UNITS/GM

A060734 001

BACITRACIN

LILLY

500 UNITS/GM

A060687 001

PHARMADERM

500 UNITS/GM

A062158 001

PHARMAFAIR

500 UNITS/GM

A062453 001 Mar 28, 1984

OINTMENT; TOPICAL

BACITRACIN

COMBE

500 UNITS/GM

A062799 001 May 14, 1987

NASKA

500 UNITS/GM

A062857 001 Nov 13, 1987

POWDER; FOR RX COMPOUNDING

BACI-RX

X GEN PHARMS

5,000,000 UNITS/BOT

A061580 001

BACITRACIN

APOTHEKERNES

5,000,000 UNITS/BOT

A061699 001

PADDOCK LLC

5,000,000 UNITS/BOT

A062456 001 Jul 27, 1983

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

BACITRACIN ZINC

POWDER;FOR RX COMPOUNDING

ZIBA-RX

X GEN PHARMS 500,000 UNITS/BOT A061737 001

BACITRACIN ZINC; HYDROCORTISONE; NEOMYCIN SULFATE; POLYMYXIN B SULFATE

OINTMENT;OPHTHALMIC

CORTISPORIN

+ CASPER PHARMA LLC 400 UNITS/GM;1%;EQ 3.5MG BASE/GM;10,000 UNITS/GM ** N050416 002

NEOMYCIN AND POLYMYXIN B SULFATES, BACITRACIN ZINC AND HYDROCORTISONE

SCIEGEN PHARMS INC 400 UNITS/GM;1%;EQ 3.5MG BASE/GM;10,000 UNITS/GM A065213 001 Jul 25, 2012

ZINC BACITRACIN,NEOMYCIN SULFATE,POLYMYXIN B SULFATE & HYDROCORTISONE

PHARMAFAIR 400 UNITS/GM;1%;EQ 3.5MG BASE/GM;10,000 UNITS/GM A062389 001 Jul 02, 1982

OINTMENT;TOPICAL

CORTISPORIN

+ MONARCH PHARMS 400 UNITS/GM;1%;EQ 3.5MG BASE/GM;5,000 UNITS/GM ** N050168 002 May 04, 1984

NEOMYCIN & POLYMYXIN B SULFATES & BACITRACIN ZINC & HYDROCORTISONE

PHARMAFAIR 400 UNITS/GM;1%;EQ 3.5MG BASE/GM;5,000 UNITS/GM A062381 001 Sep 06, 1985

BACITRACIN ZINC; LIDOCAINE; NEOMYCIN SULFATE; POLYMYXIN B SULFATE

OINTMENT;TOPICAL

LANABIOTIC

COMBE 400 UNITS/GM;40MG/GM;EQ 5MG BASE/GM;5,000 UNITS/GM A062499 001 Jun 03, 1985

BACITRACIN ZINC; NEOMYCIN SULFATE; POLYMYXIN B SULFATE

OINTMENT;OPHTHALMIC

BACITRACIN ZINC-NEOMYCIN SULFATE-POLYMYXIN B SULFATE

PHARMAFAIR 400 UNITS/GM;EQ 3.5MG BASE/GM;10,000 UNITS/GM A062386 001 Sep 09, 1982

BACITRACIN-NEOMYCIN-POLYMYXIN

PHARMADERM 400 UNITS/GM;EQ 3.5MG BASE/GM;5,000 UNITS/GM A062167 001

NEO-POLYCIN

DOW PHARM 500 UNITS/GM;EQ 3.5MG BASE/GM;10,000 UNITS/GM A060647 001

NEOMYCIN AND POLYMYXIN B SULFATES AND BACITRACIN ZINC

SCIEGEN PHARMS INC 400 UNITS/GM;EQ 3.5MG BASE/GM;10,000 UNITS/GM A065088 001 Feb 06, 2004

OINTMENT;TOPICAL

BACITRACIN ZINC-NEOMYCIN SULFATE-POLYMYXIN B SULFATE

NASKA 400 UNITS/GM;EQ 3.5MG BASE/GM;5,000 UNITS/GM A062833 001 Nov 09, 1987

BACITRACIN ZINC; POLYMYXIN B SULFATE

AEROSOL;TOPICAL

POLYSPORIN

GLAXOSMITHKLINE 10,000 UNITS/GM;2,000,000 UNITS/GM N050167 002 Mar 01, 1985

OINTMENT;OPHTHALMIC

BACITRACIN ZINC AND POLYMYXIN B SULFATE

SCIEGEN PHARMS INC 500 UNITS/GM;10,000 UNITS/GM A064028 001 Jan 30, 1995

OCUMYCIN

PHARMAFAIR 500 UNITS/GM;10,000 UNITS/GM A062430 001 Apr 08, 1983

POLYSPORIN

MONARCH PHARMS 500 UNITS/GM;10,000 UNITS/GM ** A061229 001

OINTMENT;TOPICAL

BACITRACIN ZINC-POLYMYXIN B SULFATE

NASKA 500 UNITS/GM;10,000 UNITS/GM A062849 001 Nov 13, 1987

BACITRACIN; HYDROCORTISONE ACETATE; NEOMYCIN SULFATE; POLYMYXIN B SULFATE

OINTMENT;OPHTHALMIC

BACITRACIN-NEOMYCIN-POLYMYXIN W/ HYDROCORTISONE ACETATE

ALTANA 400 UNITS/GM;1%;EQ 3.5MG BASE/GM;10,000 UNITS/GM A060731 002

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

BACITRACIN; NEOMYCIN SULFATE; POLYMYXIN B SULFATE

OINTMENT;OPHTHALMIC

MYCITRACIN

PHARMACIA AND UPJOHN	500 UNITS/GM;EQ 3.5MG BASE/GM;10,000 UNITS/GM	A061048	001	
----------------------	---	---------	-----	--

BACITRACIN; POLYMYXIN B SULFATE

DISC;TOPICAL

LANABIOTIC

COMBE	500 UNITS/GM;5,000 UNITS/GM	N050598	001	Sep 22, 1986
-------	-----------------------------	---------	-----	--------------

BACLOFEN

INJECTABLE;INTRATHECAL

BACLOFEN

HERITAGE	0.5MG/ML	A216309	001	Aug 10, 2023
	2MG/ML	A216309	002	Aug 10, 2023

SOLUTION;ORAL

OZOBAX

+ METACEL PHARMS LLC	5MG/5ML	N208193	001	Sep 18, 2019
----------------------	---------	---------	-----	--------------

TABLET;ORAL

BACLOFEN

APPCO	10MG	A090334	001	Feb 18, 2010
	20MG	A090334	002	Feb 18, 2010
IMPAX	5MG	A077971	003	Jul 07, 2021
	10MG	A077971	001	Oct 26, 2007
	20MG	A077971	002	Oct 26, 2007
NORVIUM BIOSCIENCE	10MG	A077181	001	Jul 29, 2005
	20MG	A077121	002	Jul 29, 2005
SUN PHARM INDS INC	10MG	A077862	001	Aug 14, 2006
	20MG	A077862	002	Aug 14, 2006
TEVA	10MG	A073043	001	Feb 27, 1992
	20MG	A073044	001	Feb 27, 1992
USL PHARMA	10MG	A071260	001	May 06, 1988
	20MG	A071261	001	May 06, 1988
WATSON LABS	10MG	A072824	001	Sep 18, 1991
	10MG	A073092	001	Jan 28, 1994
	10MG	A074698	001	Aug 20, 1996
	20MG	A072825	001	Sep 18, 1991
	20MG	A073093	001	Jan 28, 1994
	20MG	A074698	002	Aug 20, 1996

LIORESAL

+ NOVARTIS	10MG **	N017851	001	
+	20MG **	N017851	003	Jan 20, 1982

TABLET, ORALLY DISINTEGRATING;ORAL

KEMSTRO

+ UCB INC	10MG **	N021589	001	Oct 30, 2003
+	20MG **	N021589	002	Oct 30, 2003

BALOXAVIR MARBOXIL

TABLET;ORAL

XOFLUZA

+ GENENTECH INC	20MG **	N210854	001	Oct 24, 2018
-----------------	---------	---------	-----	--------------

BALSALAZIDE DISODIUM

CAPSULE;ORAL

BALSALAZIDE DISODIUM

ESJAY PHARMA	750MG	A077807	001	Dec 28, 2007
--------------	-------	---------	-----	--------------

TABLET;ORAL

BALSALAZIDE DISODIUM

STRIDES PHARMA	1.1GM	A206336	001	Sep 08, 2015
----------------	-------	---------	-----	--------------

GIAZO

+ VALEANT PHARMS INTL	1.1GM **	N022205	001	Feb 03, 2012
-----------------------	----------	---------	-----	--------------

BARICITINIB

TABLET;ORAL

BARICITINIB

AUROBINDO PHARMA LTD	1MG	A217542	001	Jul 22, 2024
	2MG	A217542	002	Jul 22, 2024

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

BARIUM SULFATE

FOR SUSPENSION;ORAL

E-Z-CAT DRY

+ BRACCO 40% (9GM/POUCH) N208036 003 Jan 03, 2017

BECLOMETHASONE DIPROPIONATE

AEROSOL, METERED; INHALATION

BECLOVENT

GLAXOSMITHKLINE 0.042MG/INH N018153 001

QVAR 40

+ TEVA BRANDED PHARM 0.04MG/INH ** N020911 002 Sep 15, 2000

QVAR 80

+ TEVA BRANDED PHARM 0.08MG/INH ** N020911 001 Sep 15, 2000

VANCERIL

SCHERING 0.042MG/INH N017573 001

VANCERIL DOUBLE STRENGTH

SCHERING 0.084MG/INH N020486 001 Dec 24, 1996

AEROSOL, METERED;NASAL

BECONASE

GLAXOSMITHKLINE 0.042MG/INH N018584 001

VANCENASE

SCHERING 0.042MG/INH N018521 001

BECLOMETHASONE DIPROPIONATE MONOHYDRATE

SPRAY, METERED;NASAL

BECONASE AQ

+ GLAXOSMITHKLINE EQ 0.042MG DIPROP/SPRAY N019389 001 Jul 27, 1987

VANCENASE AQ

SCHERING EQ 0.042MG DIPROP/SPRAY N019589 001 Dec 23, 1987

EQ 0.084MG DIPROP/SPRAY N020469 001 Jun 26, 1996

BENAZEPRIL HYDROCHLORIDE

TABLET;ORAL

BENAZEPRIL HYDROCHLORIDE

GENPHARM 5MG A076476 001 Feb 11, 2004

10MG A076476 002 Feb 11, 2004

20MG A076476 003 Feb 11, 2004

40MG A076476 004 Feb 11, 2004

RISING 5MG A076430 001 Feb 11, 2004

10MG A076430 002 Feb 11, 2004

20MG A076430 003 Feb 11, 2004

40MG A076430 004 Feb 11, 2004

LOTENSIN

+ VALIDUS PHARMS 5MG ** N019851 001 Jun 25, 1991

BENAZEPRIL HYDROCHLORIDE; HYDROCHLOROTHIAZIDE

TABLET;ORAL

BENAZEPRIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE

ANI PHARMS 5MG;6.25MG A076348 001 Feb 11, 2004

10MG;12.5MG A076348 002 Feb 11, 2004

20MG;12.5MG A076348 003 Feb 11, 2004

20MG;25MG A076348 004 Feb 11, 2004

AUROBINDO PHARMA USA 5MG;6.25MG A076688 001 Feb 11, 2004

10MG;12.5MG A076688 002 Feb 11, 2004

20MG;12.5MG A076688 003 Feb 11, 2004

20MG;25MG A076688 004 Feb 11, 2004

MYLAN PHARMS INC 5MG;6.25MG A076612 001 Feb 11, 2004

10MG;12.5MG A076612 002 Feb 11, 2004

20MG;12.5MG A076612 003 Feb 11, 2004

20MG;25MG A076612 004 Feb 11, 2004

SUN PHARM INDS LTD 5MG;6.25MG A077483 001 Sep 08, 2005

10MG;12.5MG A077483 002 Sep 08, 2005

20MG;12.5MG A077483 003 Sep 08, 2005

20MG;25MG A077483 004 Sep 08, 2005

LOTENSIN HCT

+ VALIDUS PHARMS 5MG;6.25MG ** N020033 001 May 19, 1992

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

BENDAMUSTINE HYDROCHLORIDE

POWDER; INTRAVENOUS

BENDAMUSTINE HYDROCHLORIDE

BRECKENRIDGE	25MG/VIAL	A205447 001	Feb 14, 2023
	100MG/VIAL	A205447 002	Feb 14, 2023
NANG KUANG PHARM CO	25MG/VIAL	A206554 001	Jun 07, 2023
	100MG/VIAL	A206554 002	Jun 07, 2023
NORVIUM BIOSCIENCE	25MG/VIAL	A204104 001	Apr 28, 2023
	100MG/VIAL	A204104 002	Apr 28, 2023

SOLUTION; INTRAVENOUS

BENDAMUSTINE HYDROCHLORIDE

+ HOSPIRA	25MG/ML (25MG/ML)	N211530 001	Dec 15, 2022
+	100MG/4ML (25MG/ML)	N211530 002	Dec 15, 2022
+	200MG/8ML (25MG/ML)	N211530 003	Dec 15, 2022
TREANDA			
+ CEPHALON	45MG/0.5ML (90MG/ML)	N022249 003	Sep 13, 2013
+	180MG/2ML (90MG/ML)	N022249 004	Sep 13, 2013

BENDROFLUMETHIAZIDE

TABLET; ORAL

NATURETIN-10

APOTHECON	10MG	N012164 003	
-----------	------	-------------	--

NATURETIN-2.5

APOTHECON	2.5MG	N012164 001	
-----------	-------	-------------	--

NATURETIN-5

APOTHECON	5MG	N012164 002	
-----------	-----	-------------	--

BENDROFLUMETHIAZIDE; NADOLOL

TABLET; ORAL

CORZIDE

+ KING PHARMS LLC	5MG;40MG **	N018647 001	May 25, 1983
+	5MG;80MG **	N018647 002	May 25, 1983

NADOLOL AND BENDROFLUMETHIAZIDE

IMPAX LABS	5MG;40MG	A077833 001	Mar 30, 2007
	5MG;80MG	A077833 002	Mar 30, 2007
NATCO PHARMA	5MG;40MG	A078688 001	Feb 15, 2008
	5MG;80MG	A078688 002	Feb 15, 2008

BENOXINATE HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

BENOXINATE HYDROCHLORIDE

SOLA BARNES HIND	0.4%	A084149 001	
------------------	------	-------------	--

BENTIROMIDE

SOLUTION; ORAL

CHYMEX

SAVAGE LABS	500MG/7.5ML	N018366 001	Dec 29, 1983
-------------	-------------	-------------	--------------

BENZONATATE

CAPSULE; ORAL

BENZONATATE

ACELLA	100MG	A091310 001	Jan 16, 2015
	200MG	A091310 002	Jan 16, 2015
ASCENT PHARMS INC	150MG	A211518 002	Feb 22, 2019
CHARTWELL RX	100MG	A210562 001	Nov 09, 2018
	150MG	A210562 002	Nov 09, 2018
	200MG	A210562 003	Nov 09, 2018
MIKART	100MG	A040851 001	Nov 09, 2009
	150MG	A040851 002	Nov 09, 2009
	200MG	A040851 003	Nov 09, 2009
NESHER PHARMS	100MG	A040795 001	Oct 31, 2007
	200MG	A040795 002	Oct 31, 2007
SUN PHARM INDS INC	100MG	A040587 001	Mar 19, 2008
	200MG	A040587 002	Mar 19, 2008
TESSALON			
+ PFIZER	200MG **	N011210 003	Jun 25, 1999

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

BENZOYL PEROXIDE; CLINDAMYCIN PHOSPHATE

GEL; TOPICAL

BENZACLIN

+	BAUSCH	5%;EQ 1% BASE	N050756 001	Dec 21, 2000
		5%;EQ 1% BASE	N050756 002	Apr 20, 2007

CLINDAMYCIN PHOSPHATE AND BENZOYL PEROXIDE

	CHARTWELL RX	5%;1.2%	A203688 001	Aug 25, 2016
	ENCUBE	5%;EQ 1% BASE	A204087 001	Jun 27, 2017

BENZOYL PEROXIDE; ERYTHROMYCIN

GEL; TOPICAL

AKTIPAK

+	BIOFRONTERA	5%;3%	N050769 001	Nov 27, 2000
---	-------------	-------	-------------	--------------

ERYTHROMYCIN AND BENZOYL PEROXIDE

	ENCUBE	5%;3%	A065112 001	Mar 29, 2004
--	--------	-------	-------------	--------------

BENZPHETAMINE HYDROCHLORIDE

TABLET; ORAL

BENZPHETAMINE HYDROCHLORIDE

	AVET LIFESCIENCES	50MG	A202061 001	Jan 27, 2012
	EPIC PHARMA LLC	50MG	A040714 001	Oct 29, 2007
	IMPAX LABS	50MG	A040845 001	Nov 18, 2008
	SCINOPHARM TAIWAN	50MG	A040578 001	Apr 17, 2006
	SPECGX LLC	50MG	A040773 001	Apr 25, 2007
	TEDOR PHARM	25MG	A040747 002	Nov 20, 2015
		50MG	A040747 001	Mar 30, 2007

DIDREX

+	PFIZER	25MG **	N012427 003	
+		50MG **	N012427 002	

BENZQUINAMIDE HYDROCHLORIDE

INJECTABLE; INJECTION

EMETE-CON

	PFIZER	EQ 50MG BASE/VIAL	N016820 001	
--	--------	-------------------	-------------	--

SUPPOSITORY; RECTAL

EMETE-CON

	ROERIG	EQ 100MG BASE	N016818 006	
--	--------	---------------	-------------	--

BENZTHIAZIDE

TABLET; ORAL

AQUATAG

	SOLVAY	25MG	N016001 001	
		50MG	N016001 002	

BENZTHIAZIDE

	PVT FORM	50MG	A083206 001	
--	----------	------	-------------	--

EXNA

	AH ROBINS INC	50MG	N012489 001	
--	---------------	------	-------------	--

FOVANE

	PFIZER	50MG	N012128 002	
--	--------	------	-------------	--

URESE

	PFIZER	25MG	N012128 003	
--	--------	------	-------------	--

BENZTROPINE MESYLATE

INJECTABLE; INJECTION

BENZTROPINE MESYLATE

	LUITPOLD	1MG/ML	A091152 001	Mar 29, 2010
--	----------	--------	-------------	--------------

COGENTIN

+	EPIC PHARMA LLC	1MG/ML **	N012015 001	
---	-----------------	-----------	-------------	--

TABLET; ORAL

BENZTROPINE MESYLATE

	INVAGEN PHARMS	0.5MG	A090294 001	Mar 29, 2010
		1MG	A090294 002	Mar 29, 2010
		2MG	A090294 003	Mar 29, 2010
	LANNETT CO INC	0.5MG **	A088877 001	Apr 11, 1985
		1MG **	A088894 001	Apr 11, 1985
		2MG **	A088895 001	Apr 11, 1985
	OXFORD PHARMS	0.5MG	A040706 002	Feb 14, 2008
		1MG	A040706 003	Feb 14, 2008
		2MG	A040706 001	Feb 14, 2008
	QUANTUM PHARMICS	0.5MG	A088514 001	Jan 31, 1984
		1MG	A088510 001	Jan 31, 1984
		2MG	A088511 001	Jan 31, 1984

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

BENZTROPINE MESYLATE

TABLET; ORAL

BENZTROPINE MESYLATE

USL PHARMA

0.5MG

A089211 001 Jun 14, 1988

1MG

A089212 001 Jun 14, 1988

2MG

A089213 001 Jun 14, 1988

COGENTIN

+ MERCK

0.5MG **

N009193 004

+

1MG **

N009193 003

+

2MG **

N009193 002

BENZYL ALCOHOL

LOTION; TOPICAL

ULESFIA

+ SHIONOGI INC

5% **

N022129 001 Apr 09, 2009

BENZYL BENZOATE

EMULSION; TOPICAL

BENZYL BENZOATE

LANNETT

50%

A084535 001

BEPRIDIL HYDROCHLORIDE

TABLET; ORAL

BEPADIN

MEDPOINTE PHARM HLC

200MG

N019001 001 Dec 28, 1990

300MG

N019001 002 Dec 28, 1990

400MG

N019001 003 Dec 28, 1990

VASCOR

JOHNSON AND JOHNSON

200MG

N019002 001 Dec 28, 1990

300MG

N019002 002 Dec 28, 1990

400MG

N019002 003 Dec 28, 1990

BETA CAROTENE

CAPSULE; ORAL

SOLATENE

ROCHE

30MG

N017589 001

BETAMETHASONE

CREAM; TOPICAL

CELESTONE

SCHERING

0.2%

N014762 001

SYRUP; ORAL

CELESTONE

MERCK SHARP DOHME

0.6MG/5ML

N014215 002

TABLET; ORAL

CELESTONE

SCHERING

0.6MG

N012657 003

BETAMETHASONE BENZOATE

CREAM; TOPICAL

UTICORT

PARKE DAVIS

0.025%

N016998 002

GEL; TOPICAL

UTICORT

PARKE DAVIS

0.025%

N017244 001

LOTION; TOPICAL

UTICORT

PARKE DAVIS

0.025%

N017528 001

OINTMENT; TOPICAL

UTICORT

PARKE DAVIS

0.025%

N018089 001

BETAMETHASONE DIPROPIONATE

CREAM; TOPICAL

ALPHATREX

SAVAGE LABS

EQ 0.05% BASE

N019138 001 Jun 26, 1984

BETAMETHASONE DIPROPIONATE

PERRIGO NEW YORK

EQ 0.05% BASE

A072536 001 Jan 31, 1990

EQ 0.05% BASE

A074579 001 Nov 26, 1997

PHARMADERM

EQ 0.05% BASE

N019136 001 Jun 26, 1984

TARO

EQ 0.05% BASE

A071143 001 Jun 17, 1987

TEVA

EQ 0.05% BASE

A071476 001 Aug 10, 1987

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

BETAMETHASONE DIPROPIONATE

CREAM;TOPICAL				
DIPROSONE				
	SCHERING	EQ 0.05% BASE	N017536	001
CREAM, AUGMENTED;TOPICAL				
BETAMETHASONE DIPROPIONATE				
	ANDA REPOSITORY	EQ 0.05% BASE	A076603	001 Jan 23, 2004
DIPROLENE				
	SCHERING	EQ 0.05% BASE	N019408	001 Jan 31, 1986
DIPROLENE AF				
	+ ORGANON	EQ 0.05% BASE **	N019555	001 Apr 27, 1987
DISC;TOPICAL				
DIPROSONE				
	SCHERING	EQ 0.1% BASE	N017829	001
GEL, AUGMENTED;TOPICAL				
DIPROLENE				
	SCHERING	EQ 0.05% BASE	N019408	002 Nov 22, 1991
LOTION;TOPICAL				
ALPHATREX				
	SAVAGE LABS	EQ 0.05% BASE	A070273	001 Aug 12, 1985
BETAMETHASONE DIPROPIONATE				
	ACTAVIS MID ATLANTIC	EQ 0.05% BASE	A070281	001 Jul 31, 1985
	ALPHARMA US PHARMS	EQ 0.05% BASE	A071085	001 Feb 03, 1987
	COSETTE	EQ 0.05% BASE	A071882	001 Jun 06, 1988
	HIKMA	EQ 0.05% BASE	A209896	001 Feb 06, 2018
	SHREE HARI INTL	EQ 0.05% BASE	A070274	001 Aug 12, 1985
	TARO	EQ 0.05% BASE	A072276	001 Aug 24, 1988
		EQ 0.05% BASE	A074272	001 Sep 30, 1994
DIPROSONE				
	+ SCHERING	EQ 0.05% BASE **	N017781	001
LOTION, AUGMENTED;TOPICAL				
BETAMETHASONE DIPROPIONATE				
	ENCUBE	EQ 0.05% BASE	A206389	001 Feb 13, 2018
	HIKMA	EQ 0.05% BASE	A208849	001 Oct 11, 2019
DIPROLENE				
	+ ORGANON	EQ 0.05% BASE **	N019716	001 Aug 01, 1988
OINTMENT;TOPICAL				
ALPHATREX				
	SAVAGE LABS	EQ 0.05% BASE	N019143	001 Sep 04, 1984
BETAMETHASONE DIPROPIONATE				
	PERRIGO NEW YORK	EQ 0.05% BASE	A072526	001 Jan 31, 1990
	PHARMADERM	EQ 0.05% BASE	N019140	001 Sep 04, 1984
	TEVA	EQ 0.05% BASE	A071477	001 Aug 10, 1987
DIPROSONE				
	SCHERING	EQ 0.05% BASE	N017691	001
OINTMENT, AUGMENTED;TOPICAL				
BETAMETHASONE DIPROPIONATE				
	PADAGIS US	EQ 0.05% BASE	A206118	001 Nov 09, 2017
SPRAY;TOPICAL				
BETAMETHASONE DIPROPIONATE				
	TARO	EQ 0.05% BASE/SPRAY	A211722	001 Jun 17, 2020

BETAMETHASONE DIPROPIONATE; CALCIPOTRIENE

AEROSOL, FOAM;TOPICAL				
CALCIPOTRIENE AND BETAMETHASONE DIPROPIONATE				
	GLENMARK PHARMS LTD	0.064%;0.005%	A214688	001 Mar 21, 2023
OINTMENT;TOPICAL				
CALCIPOTRIENE AND BETAMETHASONE DIPROPIONATE				
	CHARTWELL RX	0.064%;0.005%	A201615	001 Jan 14, 2013

BETAMETHASONE DIPROPIONATE; CLOTRIMAZOLE

CREAM;TOPICAL				
LOTRISONE				
	+ ORGANON	EQ 0.05% BASE;1% **	N018827	001 Jul 10, 1984
LOTION;TOPICAL				
LOTRISONE				
	+ MERCK SHARP DOHME	EQ 0.05% BASE;1% **	N020010	001 Dec 08, 2000

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

BETAMETHASONE SODIUM PHOSPHATE

INJECTABLE; INJECTION

BETAMETHASONE SODIUM PHOSPHATE

WATSON LABS	EQ 3MG BASE/ML	A085738	001	
CELESTONE				
+ SCHERING	EQ 3MG BASE/ML **	N017561	001	

BETAMETHASONE VALERATE

AEROSOL, FOAM; TOPICAL

BETAMETHASONE VALERATE

NOVAST LABS	0.12%	A207144	001	May 24, 2017
LUXIQ				
+ NORVIUM BIOSCIENCE	0.12%	N020934	001	Feb 28, 1999

CREAM; TOPICAL

BETADERM

ROACO	EQ 0.1% BASE	N018839	001	Jun 30, 1983
BETAMETHASONE VALERATE				
PERRIGO NEW YORK	EQ 0.1% BASE	A070053	001	Jun 10, 1986
PHARMADERM	EQ 0.1% BASE	N018860	002	Aug 31, 1983
PHARMAFAIR	EQ 0.1% BASE	A070485	001	May 29, 1987
TARO	EQ 0.1% BASE	A070062	001	May 14, 1985
BETATREX				
SAVAGE LABS	EQ 0.1% BASE	N018862	001	Aug 31, 1983
VALISONE				
SCHERING	EQ 0.01% BASE	N016322	002	
	EQ 0.1% BASE	N016322	001	

LOTION; TOPICAL

BETA-VAL

COSETTE	EQ 0.1% BASE	A070072	001	Jun 27, 1985
BETAMETHASONE VALERATE				
PHARMADERM	EQ 0.1% BASE	N018870	001	Aug 31, 1983
PHARMAFAIR	EQ 0.1% BASE	A070484	001	May 29, 1987
TEVA PHARMS	EQ 0.1% BASE	A071883	001	Apr 22, 1988
BETATREX				
SAVAGE LABS	EQ 0.1% BASE	N018867	001	Aug 31, 1983
VALISONE				
SCHERING	EQ 0.1% BASE	N016932	001	

OINTMENT; TOPICAL

BETAMETHASONE VALERATE

PERRIGO NEW YORK	EQ 0.1% BASE	A071478	001	Dec 23, 1987
PHARMADERM	EQ 0.1% BASE	N018864	001	Aug 31, 1983
PHARMAFAIR	EQ 0.1% BASE	A070486	001	May 29, 1987
BETATREX				
SAVAGE LABS	EQ 0.1% BASE	N018863	001	Aug 31, 1983
VALISONE				
SCHERING	EQ 0.1% BASE	N016740	001	

BETAXOLOL HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

BETAXOLOL HYDROCHLORIDE

APOTEX INC	EQ 0.5% BASE	A075446	001	Sep 28, 2000
SCIEGEN PHARMS INC	EQ 0.5% BASE	A075386	001	Jun 30, 2000

TABLET; ORAL

KERLONE

SANOFI AVENTIS US	10MG **	N019507	001	Oct 27, 1989
	20MG **	N019507	002	Oct 27, 1989

BETAXOLOL HYDROCHLORIDE; CHLORTHALIDONE

TABLET; ORAL

KERLEDEX

SANOFI AVENTIS US	5MG;12.5MG	N019807	001	Oct 30, 1992
	10MG;12.5MG	N019807	002	Oct 30, 1992

BETAXOLOL HYDROCHLORIDE; PILOCARPINE HYDROCHLORIDE

SUSPENSION/DROPS; OPHTHALMIC

BETOPTIC PILO

ALCON	EQ 0.25% BASE;1.75%	N020619	001	Apr 17, 1997
-------	---------------------	---------	-----	--------------

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

BETAZOLE HYDROCHLORIDE

INJECTABLE; INJECTION
HISTALOG

LILLY	50MG/ML	N009344	001
-------	---------	---------	-----

BETHANECHOL CHLORIDE

INJECTABLE; INJECTION
URECHOLINE

+ TEVA BRANDED PHARM	5MG/ML **	N006536	001
----------------------	-----------	---------	-----

TABLET; ORAL

BETHANECHOL CHLORIDE

ABLE	5MG	A040492	001	Jul 27, 2004
	10MG	A040483	001	Jul 27, 2004
	25MG	A040485	001	Jul 27, 2004
	50MG	A040509	001	Jul 27, 2004
ACTAVIS ELIZABETH	5MG	A040552	001	Oct 28, 2004
	10MG	A040553	001	Oct 28, 2004
	25MG	A040554	001	Oct 28, 2004
	50MG	A040551	001	Oct 28, 2004
ASCOT	10MG	A088288	001	Jun 08, 1983
	25MG	A088289	001	Jun 08, 1983
HERITAGE PHARMA	5MG	A091256	001	May 04, 2010
	10MG	A091256	002	May 04, 2010
	25MG	A091256	003	May 04, 2010
	50MG	A091256	004	May 04, 2010
IMPAX LABS	5MG	A040721	001	Nov 01, 2006
	10MG	A040721	002	Nov 01, 2006
	25MG	A040721	003	Nov 01, 2006
	50MG	A040721	004	Nov 01, 2006
IVAX SUB TEVA PHARMS	25MG	A084689	001	
LANNETT	5MG	A084702	001	
	10MG	A084712	001	
	25MG	A084074	001	
LANNETT CO INC	5MG	A040677	002	Mar 27, 2008
	10MG	A040677	003	Mar 27, 2008
	25MG	A040677	004	Mar 27, 2008
	50MG	A040677	001	Mar 27, 2008
SANDOZ	5MG	A084353	001	
	10MG	A084378	001	
	10MG	A084379	001	
	25MG	A084383	001	
	25MG	A084384	001	
SUN PHARM INDS INC	5MG	A040897	001	Apr 22, 2009
	10MG	A040897	002	Apr 22, 2009
	25MG	A040897	003	Apr 22, 2009
	50MG	A040897	004	Apr 22, 2009
WATSON LABS	5MG	A084402	001	
	5MG	A085230	002	
	5MG	A085841	001	
	10MG	A084408	001	
	10MG	A085228	001	
	10MG	A085842	001	
	25MG	A084441	001	
	25MG	A085229	001	
	25MG	A085839	001	
	50MG	A087397	001	
	50MG	A087444	001	
WOCKHARDT	5MG	A040532	001	Sep 29, 2003
	10MG	A040533	001	Sep 29, 2003
	25MG	A040534	001	Sep 29, 2003
	50MG	A040518	001	Sep 29, 2003
MYOTONACHOL				
GLENWOOD	5MG	A084188	001	
	10MG	A084188	003	
	25MG	A084188	004	
URECHOLINE				
ODYSSEY PHARMS	5MG	A089095	001	Dec 19, 1985
	10MG	A088440	001	May 29, 1984
	25MG	A088441	001	May 29, 1984
	50MG	A089096	001	Dec 19, 1985

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

BETHANECHOL CHLORIDE

TABLET; ORAL

URECHOLINE

+	TEVA BRANDED PHARM	5MG **	N006536 003
+		10MG **	N006536 002
+		25MG **	N006536 004
+		50MG **	N006536 005

BETHANIDINE SULFATE

TABLET; ORAL

TENATHAN

	ROBINS AH	10MG	N017675 001
		25MG	N017675 002

BETRIXABAN

CAPSULE; ORAL

BEVYXXA

+	PORTOLA PHARMS INC	40MG	N208383 001	Jun 23, 2017
+		80MG	N208383 002	Jun 23, 2017

BICALUTAMIDE

TABLET; ORAL

BICALUTAMIDE

	FRESENIUS KABI USA	50MG	A079045 001	May 13, 2010
	KUDCO IRELAND	50MG	A077995 001	Jul 06, 2009
	RISING	50MG	A079185 001	Jul 06, 2009
	ROXANE	50MG	A078285 001	Mar 24, 2011
	SYNTHON PHARMS	50MG	A077973 001	Jul 06, 2009
	TEVA	50MG	A076932 001	Jul 06, 2009

BIMATOPROST

SOLUTION/DROPS; OPHTHALMIC

BIMATOPROST

	HIKMA	0.03%	A203299 001	Nov 08, 2018
	LUMIGAN			
+	ABBVIE	0.03% **	N021275 001	Mar 16, 2001

SOLUTION/DROPS; TOPICAL

BIMATOPROST

	HIKMA	0.03%	A203051 001	Oct 09, 2018
--	-------	-------	-------------	--------------

BIPERIDEN HYDROCHLORIDE

TABLET; ORAL

AKINETON

	ABBVIE	2MG	N012003 001
--	--------	-----	-------------

BIPERIDEN LACTATE

INJECTABLE; INJECTION

AKINETON

	ABBVIE	5MG/ML	N012418 002
--	--------	--------	-------------

BISACODYL; POLYETHYLENE GLYCOL 3350; POTASSIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE

FOR SOLUTION, TABLET, DELAYED RELEASE; ORAL

HALFLYTELY

+	BRAINTREE	5MG, N/A; N/A, 210GM; N/A, 0.74GM; N/A, 2.86GM; N/A, 5.6GM **	N021551 003	Jul 16, 2010
---	-----------	---	-------------	--------------

PEG-3350, SODIUM CHLORIDE, SODIUM BICARBONATE, POTASSIUM CHLORIDE AND BISACODYL

	NOVEL LABS INC	5MG, N/A; N/A, 210GM; N/A, 0.74GM; N/A, 2.86GM; N/A, 5.6GM	A202217 001	Aug 20, 2014
--	----------------	--	-------------	--------------

BISMUTH SUBSALICYLATE; METRONIDAZOLE; TETRACYCLINE HYDROCHLORIDE

TABLET, CHEWABLE, TABLET, CAPSULE; ORAL

HELIDAC

+	CASPER PHARMA LLC	262.4MG, N/A, N/A; N/A, 250MG, N/A; N/A, N/A, 500MG **	N050719 001	Aug 15, 1996
---	-------------------	--	-------------	--------------

BISOPROLOL FUMARATE

TABLET; ORAL

BISOPROLOL FUMARATE

	DASH PHARMS NATCO	5MG	A075831 001	Dec 14, 2005
		10MG	A075831 002	Dec 14, 2005
	TEVA PHARMS	5MG	A075644 001	Jun 26, 2001
		10MG	A075644 002	Jun 26, 2001

ZEBETA

+	TEVA WOMENS	5MG **	N019982 002	Jul 31, 1992
---	-------------	--------	-------------	--------------

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

BISOPROLOL FUMARATE

TABLET; ORAL

ZEBETA

+

10MG **

N019982 001 Jul 31, 1992

BISOPROLOL FUMARATE; HYDROCHLOROTHIAZIDE

TABLET; ORAL

BISOPROLOL FUMARATE AND HYDROCHLOROTHIAZIDE

ACTAVIS ELIZABETH 2.5MG; 6.25MG

A075672 001 Sep 25, 2000

5MG; 6.25MG

A075672 002 Sep 25, 2000

10MG; 6.25MG

A075672 003 Sep 25, 2000

APOTHECON 2.5MG; 6.25MG

A075642 002 Dec 27, 2000

5MG; 6.25MG

A075642 001 Dec 27, 2000

10MG; 6.25MG

A075642 003 Dec 27, 2000

CHARTWELL RX 2.5MG; 6.25MG

A075527 001 Sep 25, 2000

5MG; 6.25MG

A075527 003 Sep 25, 2000

10MG; 6.25MG

A075527 002 Sep 25, 2000

IVAX SUB TEVA PHARMS 2.5MG; 6.25MG

A075632 001 Sep 27, 2000

5MG; 6.25MG

A075632 002 Sep 27, 2000

10MG; 6.25MG

A075632 003 Sep 27, 2000

TEVA 2.5MG; 6.25MG

A075686 001 Jan 19, 2001

5MG; 6.25MG

A075686 002 Jan 19, 2001

10MG; 6.25MG

A075686 003 Jan 19, 2001

WATSON LABS TEVA 2.5MG; 6.25MG

A075469 001 Sep 25, 2000

5MG; 6.25MG

A075469 002 Sep 25, 2000

10MG; 6.25MG

A075469 003 Sep 25, 2000

BITOLTEROL MESYLATE

AEROSOL, METERED; INHALATION

TORNALATE

SANOFI AVENTIS US 0.37MG/INH

N018770 001 Dec 28, 1984

SOLUTION; INHALATION

TORNALATE

SANOFI AVENTIS US 0.2%

N019548 001 Feb 19, 1992

BIVALIRUDIN

INJECTABLE; INTRAVENOUS

BIVALIRUDIN

APOTEX 250MG/VIAL

A204876 001 Jul 06, 2017

HOSPIRA 250MG/VIAL

A090811 001 Jul 14, 2015

250MG/VIAL

A090816 001 Jul 14, 2015

SOLUTION; INTRAVENOUS

BIVALIRUDIN IN 0.9% SODIUM CHLORIDE

+ BAXTER HLTHCARE CORP 250MG/50ML (5MG/ML)

N208374 001 Dec 21, 2017

+ 500MG/100ML (5MG/ML)

N208374 002 Dec 21, 2017

BLEOMYCIN SULFATE

INJECTABLE; INJECTION

BLENOXANE

+ BRISTOL MYERS SQUIBB EQ 15 UNITS BASE/VIAL **

N050443 001

+ EQ 30 UNITS BASE/VIAL **

N050443 002 Sep 07, 1995

BLEOMYCIN SULFATE

CIPLA EQ 15 UNITS BASE/VIAL

A209439 001 Mar 11, 2019

PHARMACHEMIE BV EQ 15 UNITS BASE/VIAL

A065201 001 Dec 13, 2007

TEVA PARENTERAL EQ 15 UNITS BASE/VIAL

A064084 001 Jun 01, 1996

EQ 30 UNITS BASE/VIAL

A064084 002 Jun 01, 1996

TEVA PHARMS USA EQ 15 UNITS BASE/VIAL

A065033 001 Jun 27, 2000

EQ 30 UNITS BASE/VIAL

A065033 002 Jun 27, 2000

BOCEPREVIR

CAPSULE; ORAL

VICTRELIS

MERCK SHARP DOHME 200MG

N202258 001 May 13, 2011

BORTEZOMIB

INJECTABLE; INTRAVENOUS, SUBCUTANEOUS

BORTEZOMIB

ACCORD HLTHCARE 3.5MG/VIAL

A204405 001 Jul 26, 2022

NORVIUM BIOSCIENCE 3.5MG/VIAL

A205160 001 Oct 31, 2022

SCINOPHARM TAIWAN 3.5MG/VIAL

A216912 001 Sep 26, 2023

TEVA PHARMS USA 3.5MG/VIAL

A205857 001 May 02, 2022

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

BORTEZOMIB

POWDER; INTRAVENOUS

BORTEZOMIB

DR REDDYS	3.5MG/VIAL	N206927	001	Oct 04,	2019
FRESENIUS KABI USA	3.5MG/VIAL	N205004	001	Nov 06,	2017

SOLUTION; INTRAVENOUS

BORTEZOMIB

+	ACCORD HLTHCARE	2.5MG/ML (2.5MG/ML) **	N215441	001	Jul 26,	2022
+		3.5MG/1.4ML (2.5MG/ML) **	N215441	002	Jul 26,	2022
+	MAIA PHARMS INC	3.5MG/3.5ML (1MG/ML)	N215331	001	Jul 27,	2022
+		3.5MG/1.4ML (2.5MG/ML)	N215331	002	Jul 27,	2022

BOSENTAN

TABLET; ORAL

BOSENTAN

ALEMBIC	62.5MG	A211461	001	Jan 23,	2020
	125MG	A211461	002	Jan 23,	2020
ALVOGEN PINE BROOK	62.5MG	A206002	001	Apr 26,	2019
	125MG	A206002	002	Apr 26,	2019
AMNEAL PHARMS CO	62.5MG	A209742	001	Apr 26,	2019
	125MG	A209742	002	Apr 26,	2019
CHARTWELL MOLECULAR	62.5MG	A210342	001	Jan 03,	2020
	125MG	A210342	002	Jan 03,	2020
ENDO OPERATIONS	62.5MG	A205699	001	Apr 26,	2019
	125MG	A205699	002	Apr 26,	2019
HIKMA	62.5MG	A208695	001	Apr 26,	2019
	125MG	A208695	002	Apr 26,	2019
MYLAN	62.5MG	A205173	001	Jan 15,	2020
	125MG	A205173	002	Jan 15,	2020
NATCO PHARMA LTD	62.5MG	A206987	001	Apr 26,	2019
	125MG	A206987	002	Apr 26,	2019

BRETYLIUM TOSYLATE

INJECTABLE; INJECTION

BRETYLIUM TOSYLATE

ABRAXIS PHARM	50MG/ML	A070134	001	Apr 29,	1986	
	100MG/ML	A071298	001	Feb 13,	1987	
ASTRAZENECA	50MG/ML	A071151	001	Aug 10,	1987	
	50MG/ML	A071152	001	Aug 10,	1987	
	50MG/ML	A071153	001	Aug 10,	1987	
BRECKENRIDGE	50MG/ML	A204386	001	Dec 21,	2018	
HIKMA	50MG/ML	A070545	001	May 14,	1986	
	50MG/ML	A070546	001	May 14,	1986	
+	HOSPIRA	50MG/ML **	N019030	001	Apr 29,	1986
	50MG/ML	N019033	001	Apr 29,	1986	
INTL MEDICATION	50MG/ML	A070119	001	Apr 29,	1986	
LUITPOLD	50MG/ML	A070891	001	Jul 26,	1988	
BRETYLIUM TOSYLATE IN DEXTROSE 5%						
ABBOTT	200MG/100ML	N019005	002	Apr 29,	1986	
	400MG/100ML	N019005	003	Apr 29,	1986	
	800MG/100ML	N019005	001	Apr 29,	1986	
BRETYLIUM TOSYLATE IN DEXTROSE 5% IN PLASTIC CONTAINER						
B BRAUN	100MG/100ML	N019121	001	Apr 29,	1986	
	200MG/100ML	N019121	002	Apr 29,	1986	
	400MG/100ML	N019121	003	Apr 29,	1986	
BAXTER HLTHCARE	200MG/100ML	N019837	002	Apr 12,	1989	
	400MG/100ML	N019837	001	Apr 12,	1989	
HOSPIRA INC	200MG/100ML	N019008	002	Apr 29,	1986	
	400MG/100ML	N019008	003	Apr 29,	1986	
	800MG/100ML	N019008	001	Apr 29,	1986	
BRETYLOL						
HOSPIRA	50MG/ML	N017954	001			

BREXPIRAZOLE

TABLET; ORAL

BREXPIRAZOLE

AJANTA PHARMA LTD	0.25MG	A213718	001	Feb 03,	2023
	0.5MG	A213718	002	Feb 03,	2023
	1MG	A213718	003	Feb 03,	2023
	2MG	A213718	004	Feb 03,	2023
	3MG	A213718	005	Feb 03,	2023

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

BREXPIRAZOLE

TABLET; ORAL

BREXPIRAZOLE

	4MG	A213718	006	Feb 03, 2023
ALKEM LABS LTD	0.25MG	A213782	001	Nov 28, 2023
	0.5MG	A213782	002	Nov 28, 2023
	1MG	A213782	003	Nov 28, 2023
	2MG	A213782	004	Nov 28, 2023
	3MG	A213782	005	Nov 28, 2023
	4MG	A213782	006	Nov 28, 2023
AMNEAL	0.25MG	A213562	001	Jan 31, 2023
	0.5MG	A213562	002	Jan 31, 2023
	1MG	A213562	003	Jan 31, 2023
	2MG	A213562	004	Jan 31, 2023
	3MG	A213562	005	Jan 31, 2023
	4MG	A213562	006	Jan 31, 2023
ANDA REPOSITORY	0.25MG	A213758	001	Oct 25, 2023
	0.5MG	A213758	002	Oct 25, 2023
	1MG	A213758	003	Oct 25, 2023
	2MG	A213758	004	Oct 25, 2023
	3MG	A213758	005	Oct 25, 2023
	4MG	A213758	006	Oct 25, 2023
AUROBINDO PHARMA LTD	0.25MG	A213659	001	Nov 02, 2023
	0.5MG	A213659	002	Nov 02, 2023
	1MG	A213659	003	Nov 02, 2023
	2MG	A213659	004	Nov 02, 2023
	3MG	A213659	005	Nov 02, 2023
	4MG	A213659	006	Nov 02, 2023
HETERO LABS LTD V	0.25MG	A213669	001	Nov 20, 2023
	0.5MG	A213669	002	Nov 20, 2023
	1MG	A213669	003	Nov 20, 2023
	2MG	A213669	004	Nov 20, 2023
	3MG	A213669	005	Nov 20, 2023
	4MG	A213669	006	Nov 20, 2023
LUPIN LTD	0.25MG	A213512	001	Mar 17, 2023
	0.5MG	A213512	002	Mar 17, 2023
	1MG	A213512	003	Mar 17, 2023
	2MG	A213512	004	Mar 17, 2023
	3MG	A213512	005	Mar 17, 2023
	4MG	A213512	006	Mar 17, 2023
SANDOZ	0.25MG	A213570	001	Sep 26, 2022
	0.5MG	A213570	002	Sep 26, 2022
	1MG	A213570	003	Sep 26, 2022
	2MG	A213570	004	Sep 26, 2022
	3MG	A213570	005	Sep 26, 2022
	4MG	A213570	006	Sep 26, 2022
TEVA PHARMS USA INC	0.25MG	A213692	001	Aug 11, 2022
	0.5MG	A213692	002	Aug 11, 2022
	1MG	A213692	003	Aug 11, 2022
	2MG	A213692	004	Aug 11, 2022
	3MG	A213692	005	Aug 11, 2022
	4MG	A213692	006	Aug 11, 2022
ZYDUS PHARMS	0.25MG	A213660	001	Jan 10, 2023
	0.5MG	A213660	002	Jan 10, 2023
	1MG	A213660	003	Jan 10, 2023
	2MG	A213660	004	Jan 10, 2023
	3MG	A213660	005	Jan 10, 2023
	4MG	A213660	006	Jan 10, 2023

BRIMONIDINE TARTRATE

SOLUTION/DROPS; OPHTHALMIC

ALPHAGAN

+ ALLERGAN	0.2% **	N020613	001	Sep 06, 1996
	0.5%	N020490	001	Mar 13, 1997

BRIMONIDINE TARTRATE

TEVA PARENTERAL	0.2%	A076372	001	Sep 10, 2004
UPSHER SMITH LABS	0.15%	A216772	001	Jun 12, 2024

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

BRIVARACETAM

TABLET; ORAL

BRIVARACETAM

AUROBINDO PHARMA LTD	50MG	A214848	001	Jan 06, 2023
	100MG	A214848	002	Jan 06, 2023
LUPIN LTD	10MG	A214918	001	Dec 20, 2022
	25MG	A214918	002	Dec 20, 2022
	50MG	A214918	003	Dec 20, 2022
	75MG	A214918	004	Dec 20, 2022
	100MG	A214918	005	Dec 20, 2022
SUNSHINE	10MG	A214748	001	Jun 09, 2022
	25MG	A214748	002	Jun 09, 2022
	50MG	A214748	003	Jun 09, 2022
	75MG	A214748	004	Jun 09, 2022
	100MG	A214748	005	Jun 09, 2022
ZYDUS PHARMS	10MG	A214501	001	Oct 03, 2022
	25MG	A214501	002	Oct 03, 2022
	50MG	A214501	003	Oct 03, 2022
	75MG	A214501	004	Oct 03, 2022
	100MG	A214501	005	Oct 03, 2022

BROMFENAC SODIUM

SOLUTION/DROPS; OPHTHALMIC

BROMDAY

+ BAUSCH AND LOMB INC	EQ 0.09% ACID **	N021664	002	Oct 16, 2010
-----------------------	------------------	---------	-----	--------------

BROMFENAC SODIUM

AMRING PHARMS	EQ 0.09% ACID	A202030	001	Jan 09, 2013
APOTEX	EQ 0.09% ACID	A202435	001	Jun 19, 2014
	EQ 0.09% ACID	A202620	001	Jun 23, 2014
CHARTWELL RX	EQ 0.09% ACID	A201941	001	Feb 10, 2015
COASTAL PHARMS	EQ 0.09% ACID	A201211	001	May 11, 2011
RISING	EQ 0.09% ACID	A203368	001	Jun 03, 2019

XIBROM

+ BAUSCH AND LOMB INC	EQ 0.09% ACID **	N021664	001	Mar 24, 2005
-----------------------	------------------	---------	-----	--------------

BROMOCRIPTINE MESYLATE

CAPSULE; ORAL

BROMOCRIPTINE MESYLATE

LEK PHARM	EQ 5MG BASE	A075100	001	Dec 10, 1998
-----------	-------------	---------	-----	--------------

TABLET; ORAL

BROMOCRIPTINE MESYLATE

AUROBINDO PHARMA USA	EQ 2.5MG BASE	A076962	001	Sep 24, 2004
----------------------	---------------	---------	-----	--------------

BROMODIPHENHYDRAMINE HYDROCHLORIDE

CAPSULE; ORAL

AMBODRYL

PARKE DAVIS	25MG	N007984	001	
-------------	------	---------	-----	--

BROMODIPHENHYDRAMINE HYDROCHLORIDE; CODEINE PHOSPHATE

SYRUP; ORAL

AMBENYL

FOREST LABS	12.5MG/5ML; 10MG/5ML	N009319	006	Jan 10, 1984
-------------	----------------------	---------	-----	--------------

BROMANYL

ALPHARMA US PHARMS	12.5MG/5ML; 10MG/5ML	A088343	001	Aug 15, 1984
--------------------	----------------------	---------	-----	--------------

BROMODIPHENHYDRAMINE HYDROCHLORIDE AND CODEINE PHOSPHATE

WOCKHARDT	12.5MG/5ML; 10MG/5ML	A088626	001	Oct 12, 1984
-----------	----------------------	---------	-----	--------------

BROMPHENIRAMINE MALEATE

ELIXIR; ORAL

BROMPHENIRAMINE MALEATE

ALPHARMA US PHARMS	2MG/5ML	A086936	001	
KV PHARM	2MG/5ML	A085466	001	
PHARM ASSOC	2MG/5ML	A087517	001	
USL PHARMA	2MG/5ML	A087964	001	Jan 25, 1983

INJECTABLE; INJECTION

BROMPHENIRAMINE MALEATE

WATSON LABS	10MG/ML	A083821	001	
	100MG/ML	A083820	001	

DIMETANE-TEN

WYETH AYERST	10MG/ML	N011418	002	
--------------	---------	---------	-----	--

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

BROMPHENIRAMINE MALEATE

TABLET; ORAL

BROMPHENIRAMINE MALEATE

BARR	4MG	A084468	001	
IVAX SUB TEVA PHARMS	4MG	A084351	001	
NEWTRON PHARMS	4MG	A086987	001	
NEXGEN PHARMA INC	4MG	A086187	001	
PAR PHARM	4MG	A087009	001	
PIONEER PHARMS	4MG	A088604	001	Jul 13, 1984
UPSHER SMITH LABS	4MG	A083215	001	
VITARINE	4MG	A085850	001	
WATSON LABS	4MG	A083123	001	
	4MG	A085769	001	

DIMETANE

WYETH CONS	4MG	N010799	003	
------------	-----	---------	-----	--

TABLET, EXTENDED RELEASE; ORAL

DIMETANE

WYETH CONS	8MG	N010799	010	Jun 10, 1983
	12MG	N010799	011	Jun 10, 1983

BROMPHENIRAMINE MALEATE; DEXTROMETHORPHAN HYDROBROMIDE; PSEUDOEPHEDRINE HYDROCHLORIDE

SYRUP; ORAL

BROMANATE DM

ALPHARMA US PHARMS	2MG/5ML; 10MG/5ML; 30MG/5ML	A088722	001	Mar 07, 1985
--------------------	-----------------------------	---------	-----	--------------

BROMFED-DM

WOCKHARDT	2MG/5ML; 10MG/5ML; 30MG/5ML	A089681	001	Dec 22, 1988
-----------	-----------------------------	---------	-----	--------------

BROMPHENIRAMINE MALEATE, PSEUDOEPHEDRINE HYDROCHLORIDE AND DEXTROMETHORPHAN HYDROBROMIDE

BIONPHARMA	2MG/5ML; 10MG/5ML; 30MG/5ML	A203997	001	Sep 30, 2020
------------	-----------------------------	---------	-----	--------------

RHODES PHARMS	2MG/5ML; 10MG/5ML; 30MG/5ML	A202955	001	Nov 05, 2020
---------------	-----------------------------	---------	-----	--------------

DIMETANE-DX

+ ROBINS AH	2MG/5ML; 10MG/5ML; 30MG/5ML **	N019279	001	Aug 24, 1984
-------------	--------------------------------	---------	-----	--------------

BROMPHENIRAMINE MALEATE; PSEUDOEPHEDRINE HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL

EFIDAC 24 PSEUDOEPHEDRINE HYDROCHLORIDE/BROMPHENIRAMINE MALEATE

ALZA	16MG; 240MG	N019672	001	Mar 29, 1996
------	-------------	---------	-----	--------------

BUCLIZINE HYDROCHLORIDE

TABLET; ORAL

BUCLADIN-S

STUART PHARMS	50MG	N010911	006	
---------------	------	---------	-----	--

BUDESONIDE

AEROSOL, METERED; NASAL

RHINOCORT

ASTRAZENECA	0.032MG/INH	N020233	001	Feb 14, 1994
-------------	-------------	---------	-----	--------------

CAPSULE, DELAYED RELEASE; ORAL

BUDESONIDE

BARR LABS DIV TEVA	3MG	A090379	001	Apr 02, 2014
--------------------	-----	---------	-----	--------------

NATCO	3MG	A206724	001	Nov 23, 2016
-------	-----	---------	-----	--------------

SCIECURE PHARMA INC	3MG	A209041	001	Sep 28, 2017
---------------------	-----	---------	-----	--------------

ORTIKOS

+ SUN PHARM INDS INC	6MG	N211929	001	Jun 13, 2019
----------------------	-----	---------	-----	--------------

+	9MG	N211929	002	Jun 13, 2019
---	-----	---------	-----	--------------

POWDER, METERED; INHALATION

PULMICORT

ASTRAZENECA	0.16MG/INH	N020441	002	Jun 24, 1997
-------------	------------	---------	-----	--------------

	0.32MG/INH	N020441	003	Jun 24, 1997
--	------------	---------	-----	--------------

SUSPENSION; INHALATION

BUDESONIDE

EUGIA PHARMA	0.5MG/2ML	A216667	001	Nov 29, 2023
--------------	-----------	---------	-----	--------------

BUMETANIDE

INJECTABLE; INJECTION

BUMETANIDE

HOSPIRA	0.25MG/ML	A074160	001	Oct 30, 1997
---------	-----------	---------	-----	--------------

	0.25MG/ML	A074332	001	Oct 31, 1994
--	-----------	---------	-----	--------------

TEVA PARENTERAL	0.25MG/ML	A074613	001	Nov 18, 1997
-----------------	-----------	---------	-----	--------------

BUMEX

+ VALIDUS PHARMS	0.25MG/ML **	N018226	001	Feb 28, 1983
------------------	--------------	---------	-----	--------------

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

BUPIVACAINE

SOLUTION; INFILTRATION

POSIMIR

+ INNOCOLL 660MG/5ML (132MG/ML) N204803 001 Feb 01, 2021

BUPIVACAINE HYDROCHLORIDE

IMPLANT; IMPLANTATION

XARACOLL

+ INNOCOLL PHARMS 100MG N209511 001 Aug 28, 2020

INJECTABLE; INJECTION

BUPIVACAINE HYDROCHLORIDE

CIVICA

0.5% A211096 001 Feb 19, 2019

HOSPIRA

0.25% A070586 001 Mar 03, 1987

0.25% N018053 002

0.5% N018053 001

0.75% A070587 001 Mar 03, 1987

0.75% N018053 003

BUPIVACAINE HYDROCHLORIDE KIT

HOSPIRA

0.075% N019978 001 Sep 03, 1992

0.114% N019978 002 Sep 03, 1992

0.23% N019978 003 Sep 03, 1992

BUPIVACAINE HYDROCHLORIDE PRESERVATIVE FREE

INTL MEDICATED

0.25% A076012 001 Jan 09, 2002

0.5% A076012 002 Jan 09, 2002

0.75% A076012 003 Jan 09, 2002

INJECTABLE; SPINAL

BUPIVACAINE HYDROCHLORIDE

BAXTER HLTHCARE CORP

0.75% A207266 001 Jul 25, 2016

SENSORCAINE

FRESENIUS KABI USA

0.75% A071202 001 Apr 15, 1987

BUPIVACAINE HYDROCHLORIDE; EPINEPHRINE

INJECTABLE; INJECTION

BUPIVACAINE HYDROCHLORIDE AND EPINEPHRINE

HOSPIRA

0.25%; 0.005MG/ML A071166 001 Jun 16, 1988

0.5%; 0.005MG/ML A071169 001 Jun 16, 1988

0.75%; 0.005MG/ML A071171 001 Jun 16, 1988

BUPIVACAINE HYDROCHLORIDE; EPINEPHRINE BITARTRATE

INJECTABLE; INJECTION

BUPIVACAINE HYDROCHLORIDE AND EPINEPHRINE

+ HOSPIRA

0.5%; 0.0091MG/ML N022046 001 Jul 13, 1983

BUPIVACAINE HYDROCHLORIDE; LIDOCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

DUOCAINE

AMPHASTAR PHARMS INC EQ 0.375% (37.5MG/10ML); EQ 1% (100MG/10ML) N021496 001 May 23, 2003

BUPIVACAINE; MELOXICAM

SOLUTION, EXTENDED RELEASE; PERIARTICULAR

ZYNRELEF KIT

+ HERON THERAPS INC 60MG/2.3ML (29.25MG/ML); 1.8MG/2.3ML (0.88MG/ML) N211988 001 May 12, 2021

+ 300MG/10.5ML (29.25MG/ML); 9MG/10.5ML (0.88MG/ML) N211988 003 May 12, 2021

BUPRENORPHINE

FILM, EXTENDED RELEASE; TRANSDERMAL

BUPRENORPHINE

ALVOGEN

7.5MCG/HR A207490 005 Mar 21, 2023

MYLAN TECH VIATRIS

5MCG/HR A210162 001 May 03, 2021

7.5MCG/HR A210162 002 May 03, 2021

10MCG/HR A210162 003 May 03, 2021

15MCG/HR A210162 004 May 03, 2021

20MCG/HR A210162 005 May 03, 2021

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

BUPRENORPHINE HYDROCHLORIDE

FILM;BUCCAL

BUPRENORPHINE HYDROCHLORIDE

ALVOGEN	EQ 0.075MG BASE	A211594 001	Aug 03, 2021
	EQ 0.15MG BASE	A211594 002	Aug 03, 2021
	EQ 0.3MG BASE	A211594 003	Aug 03, 2021
	EQ 0.45MG BASE	A211594 004	Aug 03, 2021
	EQ 0.6MG BASE	A211594 005	Aug 03, 2021
	EQ 0.75MG BASE	A211594 006	Aug 03, 2021
	EQ 0.9MG BASE	A211594 007	Aug 03, 2021

IMPLANT;IMPLANTATION

PROBUPHINE

+ REACX PHARMS	EQ 80MG BASE/IMPLANT	N204442 001	May 26, 2016
----------------	----------------------	-------------	--------------

INJECTABLE;INJECTION

BUPRENEX

+ INDIVIOR	EQ 0.3MG BASE/ML **	N018401 001	
------------	---------------------	-------------	--

BUPRENORPHINE HYDROCHLORIDE

AM REGENT	EQ 0.3MG BASE/ML	A078331 001	Mar 27, 2007
-----------	------------------	-------------	--------------

TABLET;SUBLINGUAL

BUPRENORPHINE HYDROCHLORIDE

BARR	EQ 2MG BASE	A090360 001	May 07, 2010
	EQ 8MG BASE	A090360 002	May 07, 2010
NORVIUM BIOSCIENCE	EQ 2MG BASE	A201066 001	Mar 06, 2015
	EQ 8MG BASE	A201066 002	Mar 06, 2015

SUBUTEX

+ INDIVIOR	EQ 2MG BASE **	N020732 002	Oct 08, 2002
+	EQ 8MG BASE **	N020732 003	Oct 08, 2002

BUPRENORPHINE HYDROCHLORIDE; NALOXONE HYDROCHLORIDE

FILM;BUCCAL

BUNAVAIL

+ BDSI	EQ 2.1MG BASE;EQ 0.3MG BASE	N205637 001	Jun 06, 2014
+	EQ 4.2MG BASE;EQ 0.7MG BASE	N205637 002	Jun 06, 2014
+	EQ 6.3MG BASE;EQ 1MG BASE	N205637 003	Jun 06, 2014

FILM;SUBLINGUAL

CASSIPA

+ TEVA PHARMS USA	EQ 16MG BASE;EQ 4MG BASE	N208042 001	Sep 07, 2018
-------------------	--------------------------	-------------	--------------

TABLET;SUBLINGUAL

BUPRENORPHINE HYDROCHLORIDE AND NALOXONE HYDROCHLORIDE

DR REDDYS LABS SA	EQ 2MG BASE;EQ 0.5MG BASE	A206953 001	Jul 17, 2020
	EQ 8MG BASE;EQ 2MG BASE	A206953 002	Jul 17, 2020
TEVA PHARMS USA	EQ 2MG BASE;EQ 0.5MG BASE	A091149 001	Sep 08, 2014
	EQ 8MG BASE;EQ 2MG BASE	A091149 002	Sep 08, 2014

SUBOXONE

+ INDIVIOR	EQ 2MG BASE;EQ 0.5MG BASE **	N020733 001	Oct 08, 2002
+	EQ 8MG BASE;EQ 2MG BASE **	N020733 002	Oct 08, 2002

BUPROPION HYDROCHLORIDE

TABLET;ORAL

BUPROPION HYDROCHLORIDE

HERITAGE PHARMA	75MG	A075310 001	Nov 29, 1999
	100MG	A075310 002	Nov 29, 1999
INVAGEN PHARMS	75MG	A207389 001	Sep 18, 2017
	100MG	A207389 002	Sep 18, 2017
INVATECH	75MG	A075613 002	Oct 10, 2000
	100MG	A075613 001	Oct 10, 2000

WELLBUTRIN

+ GLAXOSMITHKLINE	50MG **	N018644 001	Dec 30, 1985
+	75MG **	N018644 002	Dec 30, 1985
+	100MG **	N018644 003	Dec 30, 1985

TABLET, EXTENDED RELEASE;ORAL

BUPROPION HYDROCHLORIDE

ACTAVIS LABS FL INC	300MG	A077715 002	Jun 13, 2007
AUROBINDO PHARMA USA	100MG	A090325 001	Apr 08, 2010
	150MG	A090325 002	Apr 08, 2010
	150MG	A090942 001	Jul 14, 2010
	200MG	A090325 003	Apr 08, 2010
	300MG	A090942 002	Jul 14, 2010
CHARTWELL RX	150MG	A076834 001	Jul 14, 2005
ENDO OPERATIONS	100MG	A091459 001	Jun 09, 2011

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

BUPROPION HYDROCHLORIDETABLET, EXTENDED RELEASE;ORAL
BUPROPION HYDROCHLORIDE

	150MG	A077284 001	Dec 14, 2006
	150MG	A091459 002	Jun 09, 2011
	150MG	A091520 001	Jun 09, 2011
	200MG	A091459 003	Jun 09, 2011
	300MG	A077284 002	Dec 14, 2006
IMPAX LABS	150MG	A077415 001	Nov 26, 2008
	200MG	A076711 001	Dec 03, 2004
	300MG	A077415 002	Dec 15, 2006
INVAGEN PHARMS	100MG	A206674 001	Feb 09, 2016
	150MG	A206556 001	Aug 26, 2016
	150MG	A206674 002	Feb 09, 2016
	200MG	A206674 003	Feb 09, 2016
	300MG	A206556 002	Aug 26, 2016
JUBILANT GENERICS	100MG	A202774 001	Oct 11, 2013
	150MG	A202774 002	Oct 11, 2013
	150MG	A202775 001	Oct 11, 2013
	150MG	A207459 001	Jun 30, 2017
	200MG	A202774 003	Oct 11, 2013
	300MG	A207459 002	Jun 30, 2017
RISING	150MG	A090941 001	May 03, 2010
SANDOZ	100MG	A076845 001	Jul 14, 2005
	150MG	A076845 002	Jul 14, 2005
SUN PHARM	100MG	A078866 001	Apr 06, 2010
	150MG	A078866 002	Apr 06, 2010
	150MG	A200695 001	Dec 18, 2014
	200MG	A078866 003	Apr 06, 2010
TORRENT	100MG	A203969 001	Oct 31, 2014
	150MG	A203969 002	Oct 31, 2014
	200MG	A203969 003	Oct 31, 2014
WATSON LABS INC	100MG	A077455 001	Jul 19, 2010
	150MG	A077455 002	Mar 12, 2008
	200MG	A077455 003	Jul 19, 2010
WOCKHARDT LTD	100MG	A201331 001	Aug 30, 2012
	150MG	A201331 002	Aug 30, 2012
	200MG	A201331 003	Aug 30, 2012
WELLBUTRIN SR			
GLAXOSMITHKLINE	50MG	N020358 001	Oct 04, 1996
ZYBAN			
GLAXOSMITHKLINE	100MG	N020711 002	May 14, 1997
+	150MG **	N020711 003	May 14, 1997

BUSPIRONE HYDROCHLORIDE

CAPSULE;ORAL

BUSPAR

+	BRISTOL MYERS SQUIBB	5MG **	N021190 001	Dec 20, 2000
+		7.5MG **	N021190 002	Dec 20, 2000
+		10MG **	N021190 003	Dec 20, 2000
+		15MG **	N021190 004	Dec 20, 2000

TABLET;ORAL

BUSPAR

+	BRISTOL MYERS SQUIBB	5MG **	N018731 001	Sep 29, 1986
+		10MG **	N018731 002	Sep 29, 1986
+		15MG **	N018731 003	Apr 22, 1996
+		30MG **	N018731 004	Apr 22, 1996

BUSPIRONE HYDROCHLORIDE

AMNEAL PHARMS CO

	5MG	A208829 001	May 24, 2017
	7.5MG	A208829 002	May 24, 2017
	10MG	A208829 003	May 24, 2017
	15MG	A208829 004	May 24, 2017
	30MG	A208829 005	May 24, 2017

EGIS

	5MG	A075119 001	Mar 14, 2002
	10MG	A075119 002	Mar 14, 2002
	15MG	A075119 003	Jan 23, 2003

IVAX SUB TEVA PHARMS

	5MG **	A075385 001	Mar 01, 2002
	10MG **	A075385 002	Mar 01, 2002
	15MG **	A075385 003	Mar 01, 2002

MYLAN

	5MG	A075467 001	Feb 28, 2002
--	-----	-------------	--------------

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

BUSPIRONE HYDROCHLORIDE

TABLET; ORAL

BUSPIRONE HYDROCHLORIDE

	7.5MG	A075467 002	Mar 28, 2001
	10MG	A075467 003	Feb 28, 2002
	15MG	A075467 004	Feb 28, 2002
NESHER PHARMS	5MG	A075572 001	Feb 27, 2002
	10MG	A075572 002	Feb 27, 2002
	15MG	A075572 003	Feb 27, 2002
RISING	5MG	A075413 001	Mar 19, 2002
	10MG	A075413 002	Mar 19, 2002
	15MG	A075413 003	Mar 19, 2002

BUSULFAN

INJECTABLE; INJECTION

BUSULFAN

ACTAVIS	6MG/ML	A205139 001	Dec 08, 2017
AM REGENT	6MG/ML	A202259 001	Dec 22, 2015
ARTHUR GRP	6MG/ML	A205106 001	Sep 21, 2018
EUGIA PHARMA	6MG/ML	A215102 001	Jun 25, 2024
NEXUS	6MG/ML	A207794 001	Jan 14, 2019
NORVIUM BIOSCIENCE	6MG/ML	A205184 001	Jul 13, 2018

BUTABARBITAL SODIUM

CAPSULE; ORAL

BUTICAPS

MEDPOINTE PHARM HLC	15MG	A085381 001	
	30MG	A085381 002	
	50MG	A085381 003	
	100MG	A085381 004	

ELIXIR; ORAL

BUTABARB

ALPHARMA US PHARMS	30MG/5ML	A085873 001	
--------------------	----------	-------------	--

BUTABARBITAL SODIUM

WOCKHARDT	30MG/5ML	A085383 001	
-----------	----------	-------------	--

BUTALAN

LANNETT	33.3MG/5ML	A085880 001	
---------	------------	-------------	--

BUTISOL SODIUM

MEDA PHARMS	30MG/5ML	A085380 001	
-------------	----------	-------------	--

SARISOL

HALSEY	30MG/5ML	A084723 001	
--------	----------	-------------	--

TABLET; ORAL

BUTABARBITAL

BUNDY	30MG	A085550 001	
-------	------	-------------	--

BUTABARBITAL SODIUM

SANDOZ	15MG	A084292 003	Feb 09, 1982
--------	------	-------------	--------------

+

	15MG	A085938 001	
--	------	-------------	--

	30MG	A084272 002	
--	------	-------------	--

	30MG	A085934 001	
--	------	-------------	--

SOLVAY	16.2MG	A083606 001	
--------	--------	-------------	--

	32.4MG	A083898 001	
--	--------	-------------	--

	48.6MG	A083897 001	
--	--------	-------------	--

	97.2MG	A083896 001	
--	--------	-------------	--

TEVA	15MG	A088632 001	May 18, 1985
------	------	-------------	--------------

	30MG	A088631 001	May 01, 1985
--	------	-------------	--------------

WATSON LABS	15MG	A085764 001	
-------------	------	-------------	--

	30MG	A085772 001	
--	------	-------------	--

WHITEWORTH TOWN PLSN	15MG	A083325 002	
----------------------	------	-------------	--

	30MG	A083337 001	
--	------	-------------	--

BUTISOL SODIUM

NORVIUM BIOSCIENCE	15MG **	N000793 002	
--------------------	---------	-------------	--

+

	30MG	N000793 004	
--	------	-------------	--

	50MG **	N000793 003	
--	---------	-------------	--

	100MG **	N000793 005	
--	----------	-------------	--

SARISOL NO. 1

HALSEY	15MG	A084719 001	
--------	------	-------------	--

SARISOL NO. 2

HALSEY	30MG	A084719 002	
--------	------	-------------	--

SODIUM BUTABARBITAL

HIKMA	15MG	A085418 001	
-------	------	-------------	--

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

BUTABARBITAL SODIUM

TABLET; ORAL

SODIUM BUTABARBITAL

	30MG	A085432	001
IVAX SUB TEVA PHARMS	15MG	A083484	001
	30MG	A084040	001
LANNETT	15MG	A085849	001
	30MG	A085866	001
	100MG	A085881	001
MARSHALL PHARMA	16.2MG	A083524	001
	32.4MG	A083858	001

BUTENAFINE HYDROCHLORIDE

CREAM; TOPICAL

MENTAX

+ NORVIUM BIOSCIENCE 1% N020524 001 Oct 18, 1996

MENTAX-TC

NORVIUM BIOSCIENCE 1% N021408 001 Oct 17, 2002

BUTOCONAZOLE NITRATE

CREAM; VAGINAL

BUTOCONAZOLE NITRATE

PADAGIS US 2% N019881 001 Feb 07, 1997

FEMSTAT

ROCHE PALO 2% N019215 001 Nov 25, 1985

FEMSTAT 3

+ BAYER 2% N020421 001 Dec 21, 1995

SUPPOSITORY; VAGINAL

FEMSTAT

ROCHE PALO 100MG N019359 001 Nov 25, 1985

BUTORPHANOL TARTRATE

INJECTABLE; INJECTION

BUTORPHANOL TARTRATE

BAXTER HLTHCARE CORP 2MG/ML A075697 001 Oct 23, 2001

HIKMA FARMACEUTICA 2MG/ML A078247 001 Apr 29, 2009

HOSPIRA 1MG/ML A075342 001 Nov 04, 1999

1MG/ML A075559 001 Mar 20, 2000

2MG/ML A075342 002 Nov 04, 1999

2MG/ML A075559 002 Mar 20, 2000

BUTORPHANOL TARTRATE PRESERVATIVE FREE

BAXTER HLTHCARE CORP 1MG/ML A075695 001 Oct 23, 2001

2MG/ML A075695 002 Oct 23, 2001

HOSPIRA 1MG/ML A074620 001 Jan 22, 1997

1MG/ML A075170 001 Sep 28, 1998

2MG/ML A074620 002 Jan 22, 1997

2MG/ML A075170 002 Sep 28, 1998

STADOL

+ APOTHECON 2MG/ML ** N017857 004

STADOL PRESERVATIVE FREE

+ APOTHECON 1MG/ML ** N017857 001

+ 2MG/ML ** N017857 002

SPRAY, METERED; NASAL

STADOL

BRISTOL MYERS SQUIBB 1MG/SPRAY ** N019890 001 Dec 12, 1991

CABAZITAXEL

SOLUTION; INTRAVENOUS

CABAZITAXEL

+ ACCORD HLTHCARE 60MG/3ML (20MG/ML) N207949 001 Dec 29, 2021

+ ACTAVIS 60MG/6ML (10MG/ML) N207970 001 Mar 14, 2024

APOTEX 60MG/1.5ML (40MG/ML) A207736 001 Feb 10, 2023

BRECKENRIDGE 60MG/1.5ML (40MG/ML) A207619 001 Jun 23, 2022

MYLAN LABS LTD 60MG/1.5ML (40MG/ML) A207381 001 Jul 05, 2023

CABERGOLINE

TABLET; ORAL

CABERGOLINE

ACTAVIS LABS FL INC 0.5MG A078035 001 Apr 21, 2008

APOTEX CORP 0.5MG A201503 001 Mar 08, 2013

IMPAX LABS INC 0.5MG A077843 001 Jul 03, 2007

NORVIUM BIOSCIENCE 0.5MG A202947 001 Dec 02, 2013

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CABERGOLINETABLET; ORAL
DOSTINEX

+ PFIZER 0.5MG ** N020664 001 Dec 23, 1996

CAFFEINE CITRATESOLUTION; INTRAVENOUS
CAFFEINE CITRATE

SUN PHARM EQ 30MG BASE/3ML (EQ 10MG BASE/ML) A090077 001 Sep 30, 2009

SOLUTION; ORAL
CAFECIT

+ HIKMA EQ 30MG BASE/3ML (EQ 10MG BASE/ML) ** N020793 002 Apr 12, 2000

CAFFEINE CITRATE
AM REGENT

EQ 30MG BASE/3ML (EQ 10MG BASE/ML) A090064 001 Nov 20, 2009

CAFFEINE; ERGOTAMINE TARTRATE

SUPPOSITORY; RECTAL

CAFERGOT

+ NOVARTIS 100MG; 2MG ** N009000 002

TABLET; ORAL

CAFERGOT

+ CHARTWELL RX 100MG; 1MG A084294 001

NOVARTIS 100MG; 1MG N006620 001

ERGOTAMINE TARTRATE AND CAFFEINE

HIKMA INTL PHARMS 100MG; 1MG A040510 001 Sep 17, 2004

WIGRAINE

ORGANON USA INC 100MG; 1MG A086562 001

CALCIFEDIOL

CAPSULE; ORAL

CALDEROL

ORGANON USA INC 0.02MG N018312 001

0.05MG N018312 002

CALCIPOTRIENE

CREAM; TOPICAL

CALCIPOTRIENE

CHARTWELL RX 0.005% A200935 001 May 30, 2012

SOLUTION; TOPICAL

CALCIPOTRIENE

CHARTWELL RX 0.005% A077029 001 Nov 20, 2009

HIKMA 0.005% A077579 001 Nov 19, 2009

DOVONEX

+ LEO PHARM 0.005% ** N020611 001 Mar 03, 1997

CALCITONIN HUMAN

INJECTABLE; INJECTION

CIBACALCIN

NOVARTIS 0.5MG/VIAL N018470 001 Oct 31, 1986

CALCITONIN SALMON

INJECTABLE; INJECTION

CALCIMAR

+ SANOFI AVENTIS US 200 IU/ML ** N017769 001

400 IU/VIAL N017497 001

CALCITONIN-SALMON

IGI LABS INC 200 IU/ML A073690 001 Apr 14, 1995

MIACALCIN

+ MYLAN IRELAND LTD 100 IU/ML ** N017808 001 Jul 03, 1986

SPRAY, METERED; NASAL

MIACALCIN

+ NORVIUM BIOSCIENCE 200 IU/SPRAY ** N020313 002 Aug 17, 1995

CALCITONIN SALMON RECOMBINANT

SPRAY, METERED; NASAL

FORTICAL

UPSHER SMITH LABS 200 IU/SPRAY ** N021406 001 Aug 12, 2005

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CALCITRIOL

CAPSULE; ORAL

CALCITRIOL

SUN PHARM	0.25MCG	A204556 001	Feb 21, 2019
	0.5MCG	A204556 002	Feb 21, 2019

INJECTABLE; INJECTION

CALCIJEX

+ ABBVIE	0.001MG/ML **	N018874 001	Sep 25, 1986
+	0.002MG/ML **	N018874 002	Sep 25, 1986

CALCITRIOL

AM REGENT	0.001MG/ML	A075746 001	Sep 26, 2003
	0.002MG/ML	A075746 002	Sep 26, 2003
FRESENIUS KABI USA	0.001MG/ML	A075836 001	Dec 31, 2002
	0.002MG/ML	A075836 002	Dec 31, 2002
FRESENIUS MEDCL	0.001MG/ML	A075766 001	Feb 20, 2003
	0.002MG/ML	A075766 002	Feb 20, 2003
HOSPIRA	0.001MG/ML	A075816 001	Jan 16, 2004
	0.002MG/ML	A075816 002	Jan 16, 2004
LONG GROVE PHARMS	0.002MG/ML	A078066 002	Jan 29, 2008
ROCKWELL MEDCL	0.001MG/ML	A076206 001	Sep 17, 2003
SAGENT PHARMS	0.001MG/ML	A077102 001	Feb 08, 2006
TEVA PARENTERAL	0.001MG/ML	A075823 001	Mar 31, 2003
	0.002MG/ML	A075823 002	Mar 31, 2003

CALCIUM ACETATE

CAPSULE; ORAL

CALCIUM ACETATE

AMNEAL PHARMS	667MG	A201658 001	Oct 06, 2014
LOTUS PHARM CO LTD	667MG	A203298 001	Jul 26, 2016

PHOSLO

FRESENIUS MEDCL	333.5MG	N021160 001	Apr 02, 2001
	667MG	N021160 002	Apr 02, 2001

PHOSLO GELCAPS

+ FRESENIUS MEDCL	667MG	N021160 003	Apr 02, 2001
-------------------	-------	-------------	--------------

SOLUTION; ORAL

PHOSLYRA

+ FRESENIUS MEDCL	667MG/5ML	N022581 001	Apr 18, 2011
-------------------	-----------	-------------	--------------

TABLET; ORAL

CALCIUM ACETATE

HIKMA	667MG	A077693 001	Jan 30, 2008
-------	-------	-------------	--------------

ELIPHOS

CYPRESS PHARM	667MG	A078502 001	Nov 25, 2008
---------------	-------	-------------	--------------

PHOSLO

+ FRESENIUS MEDCL	667MG **	N019976 001	Dec 10, 1990
-------------------	----------	-------------	--------------

CALCIUM CARBONATE; FAMOTIDINE; MAGNESIUM HYDROXIDE

TABLET, CHEWABLE; ORAL

FAMOTIDINE, CALCIUM CARBONATE, AND MAGNESIUM HYDROXIDE

PERRIGO R AND D	800MG;10MG;165MG	A204782 001	Aug 29, 2016
-----------------	------------------	-------------	--------------

CALCIUM CARBONATE; RISEDRONATE SODIUM

TABLET; ORAL

ACTONEL WITH CALCIUM (COPACKAGED)

+ WARNER CHILCOTT	EQ 500MG BASE;35MG **	N021823 001	Aug 12, 2005
-------------------	-----------------------	-------------	--------------

CALCIUM CHLORIDE; DEXTROSE; GLUTATHIONE DISULFIDE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE; SODIUM PHOSPHATE

SOLUTION; IRRIGATION

METHOTREXATE SODIUM

EPIC PHARMA LLC	0.154MG/ML;0.92MG/ML;0.184MG/ML;0.2MG/ML; L;0.38MG/ML;2.1MG/ML;7.14MG/ML;0.42MG/ML	N020079 001	Feb 26, 1999
-----------------	---	-------------	--------------

CALCIUM CHLORIDE; DEXTROSE; LACTIC ACID; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE

INJECTABLE; INJECTION

PRISMASOL B22GK 2/0 IN PLASTIC CONTAINER

+ BAXTER HLTHCARE CORP	N/A/1000ML;20GM/1000ML;5.4GM/1000ML;3.0 5GM/1000ML;0.157GM/1000ML;2.21GM/1000ML ;7.07GM/1000ML (5000ML)	N021703 010	Oct 10, 2008
------------------------	---	-------------	--------------

PRISMASOL B22GK 2/2.5 IN PLASTIC CONTAINER

BAXTER HLTHCARE CORP	3.68GM/1000ML;20GM/1000ML;5.4GM/1000ML; 3.05GM/1000ML;0.157GM/1000ML;2.21	N021703 012	Oct 10, 2008
----------------------	--	-------------	--------------

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CALCIUM CHLORIDE; DEXTROSE; LACTIC ACID; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE

INJECTABLE; INJECTION

PRISMASOL B22GK 2/2.5 IN PLASTIC CONTAINER				
		GM/1000ML; 7.07GM/1000ML (5000ML)		
PRISMASOL B22GK 4/2.5 IN PLASTIC CONTAINER				
+ BAXTER HLTHCARE CORP	3.68GM/1000ML; 20GM/1000ML; 5.4GM/1000ML; 3.05GM/1000ML; 0.314GM/1000ML; 2.21GM/1000ML; 7.07GM/1000ML (5000ML)		N021703 013	Oct 10, 2008
PRISMASOL BGK 4/0 IN PLASTIC CONTAINER				
BAXTER HLTHCARE CORP	N/A/1000ML; 20GM/1000ML; 5.4GM/1000ML; 3.05GM/1000ML; 0.314GM/1000ML; 3.09GM/1000ML; 6.46GM/1000ML (5000ML)		N021703 005	Oct 25, 2006
PRISMASOL BGK 4/3.5 IN PLASTIC CONTAINER				
BAXTER HLTHCARE CORP	5.15GM/1000ML; 20GM/1000ML; 5.4GM/1000ML; 2.03GM/1000ML; 0.314GM/1000ML; 3.09GM/1000ML; 6.46GM/1000ML (5000ML)		N021703 008	Oct 25, 2006
PRISMASOL BK 0/0 IN PLASTIC CONTAINER				
BAXTER HLTHCARE CORP	N/A/1000ML; N/A/1000ML; 5.4GM/1000ML; 3.05GM/1000ML; N/A/1000ML; 3.09GM/1000ML; 6.46GM/1000ML (5000ML)		N021703 007	Oct 25, 2006
PRISMASOL BK 0/3.5 IN PLASTIC CONTAINER				
+ BAXTER HLTHCARE CORP	5.15GM/1000ML; N/A/1000ML; 5.4GM/1000ML; 2.03GM/1000ML; N/A/1000ML; 3.09GM/1000ML; 6.46GM/1000ML (5000ML)		N021703 001	Oct 25, 2006
PRISMASOL BK 4/2.5 IN PLASTIC CONTAINER				
BAXTER HLTHCARE CORP	3.68GM/1000ML; N/A/1000ML; 5.4GM/1000ML; 3.05GM/1000ML; 0.314GM/1000ML; 3.09GM/1000ML; 6.46GM/1000ML (5000ML)		N021703 009	Oct 25, 2006

CALCIUM CHLORIDE; DEXTROSE; MAGNESIUM CHLORIDE; OXIGLUTATONE; POTASSIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE; SODIUM PHOSPHATE

SOLUTION; IRRIGATION

NAVSTEL

ALCON PHARMS LTD	0.154MG/ML; 0.92MG/ML; 0.2MG/ML; 0.184MG/ML; 0.38MG/ML; 2.1MG/ML; 7.14MG/ML; 0.42MG/ML		N022193 001	Jul 24, 2008
------------------	--	--	-------------	--------------

CALCIUM CHLORIDE; DEXTROSE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE

INJECTABLE; INJECTION

ISOLYTE R IN DEXTROSE 5% IN PLASTIC CONTAINER				
B BRAUN	37MG/100ML; 5GM/100ML; 31MG/100ML; 120MG/100ML; 330MG/100ML; 88MG/100ML		N019864 001	Jun 10, 1993
ISOLYTE R W/ DEXTROSE 5% IN PLASTIC CONTAINER				
B BRAUN	37MG/100ML; 5GM/100ML; 31MG/100ML; 120MG/100ML; 330MG/100ML; 88MG/100ML		N018271 001	

CALCIUM CHLORIDE; DEXTROSE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE; SODIUM CITRATE

INJECTABLE; INJECTION

ISOLYTE E IN DEXTROSE 5% IN PLASTIC CONTAINER				
B BRAUN	35MG/100ML; 5GM/100ML; 30MG/100ML; 74MG/100ML; 640MG/100ML; 500MG/100ML; 74MG/100ML		N019867 001	Dec 20, 1993
ISOLYTE E W/ DEXTROSE 5% IN PLASTIC CONTAINER				
B BRAUN	35MG/100ML; 5GM/100ML; 30MG/100ML; 74MG/100ML; 640MG/100ML; 500MG/100ML; 74MG/100ML		N018269 002	Jan 17, 1983

CALCIUM CHLORIDE; DEXTROSE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE; SODIUM LACTATE

INJECTABLE; INJECTION

PLASMA-LYTE M AND DEXTROSE 5% IN PLASTIC CONTAINER				
+ BAXTER HLTHCARE	37MG/100ML; 5GM/100ML; 30MG/100ML; 119MG/100ML; 161MG/100ML; 94MG/100ML; 138MG/100ML		N017390 001	
	**			

CALCIUM CHLORIDE; DEXTROSE; MAGNESIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE

SOLUTION; INTRAPERITONEAL

DIALYTE CONCENTRATE W/ DEXTROSE 30% IN PLASTIC CONTAINER				
B BRAUN	510MG/100ML; 30GM/100ML; 200MG/100ML; 9.2GM/100ML; 9.6GM/100ML		N018807 001	Aug 26, 1983
	510MG/100ML; 30GM/100ML; 200MG/100ML; 9.4GM/100ML; 11GM/100ML		N018807 003	Aug 26, 1983

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CALCIUM CHLORIDE; DEXTROSE; MAGNESIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE

SOLUTION; INTRAPERITONEAL

DIALYTE CONCENTRATE W/ DEXTROSE 50% IN PLASTIC CONTAINER

B BRAUN	510MG/100ML; 50GM/100ML; 200MG/100ML; 9.2G M/100ML; 9.6GM/100ML	N018807 002	Aug 26, 1983
---------	--	-------------	--------------

	510MG/100ML; 50GM/100ML; 200MG/100ML; 9.4G M/100ML; 11GM/100ML	N018807 004	Aug 26, 1983
--	---	-------------	--------------

DIALYTE LM/ DEXTROSE 2.5% IN PLASTIC CONTAINER

B BRAUN	29MG/100ML; 2.5GM/100ML; 15MG/100ML; 610MG /100ML; 560MG/100ML	N018460 006	Jan 29, 1986
---------	---	-------------	--------------

DIALYTE W/ DEXTROSE 1.5% IN PLASTIC CONTAINER

B BRAUN	29MG/100ML; 1.5GM/100ML; 15MG/100ML; 610MG /100ML; 560MG/100ML	N018460 001	
---------	---	-------------	--

DIALYTE W/ DEXTROSE 4.25% IN PLASTIC CONTAINER

B BRAUN	29MG/100ML; 4.25GM/100ML; 15MG/100ML; 610M G/100ML; 560MG/100ML	N018460 003	
---------	--	-------------	--

CALCIUM CHLORIDE; DEXTROSE; MAGNESIUM CHLORIDE; SODIUM CHLORIDE; SODIUM LACTATE

SOLUTION; INTRAPERITONEAL

DELFLX W/ DEXTROSE 1.5% IN PLASTIC CONTAINER

FRESENIUS MEDCL	25.7MG/100ML; 1.5GM/100ML; 15.2MG/100ML; 5 67MG/100ML; 392MG/100ML	N018379 002	
-----------------	---	-------------	--

DELFLX W/ DEXTROSE 2.5% IN PLASTIC CONTAINER

FRESENIUS MEDCL	25.7MG/100ML; 2.5GM/100ML; 15.2MG/100ML; 5 67MG/100ML; 392MG/100ML	N018379 003	
-----------------	---	-------------	--

DELFLX W/ DEXTROSE 3.5% IN PLASTIC CONTAINER

FRESENIUS MEDCL	25.7MG/100ML; 3.5GM/100ML; 15.2MG/100ML; 5 67MG/100ML; 392MG/100ML	N018379 007	Jun 24, 1988
-----------------	---	-------------	--------------

DELFLX W/ DEXTROSE 4.25% IN PLASTIC CONTAINER

FRESENIUS MEDCL	25.7MG/100ML; 4.25GM/100ML; 15.2MG/100ML; 567MG/100ML; 392MG/100ML	N018379 001	
-----------------	---	-------------	--

DELFLX-LM W/ DEXTROSE 1.5% IN PLASTIC CONTAINER

FRESENIUS MEDCL	25.7MG/100ML; 1.5GM/100ML; 5.08MG/100ML; 5 38MG/100ML; 448MG/100ML	N018379 004	Jul 07, 1982
-----------------	---	-------------	--------------

DELFLX-LM W/ DEXTROSE 2.5% IN PLASTIC CONTAINER

FRESENIUS MEDCL	25.7MG/100ML; 2.5GM/100ML; 5.08MG/100ML; 5 38MG/100ML; 448MG/100ML	N018379 005	Jul 07, 1982
-----------------	---	-------------	--------------

DELFLX-LM W/ DEXTROSE 3.5% IN PLASTIC CONTAINER

FRESENIUS MEDCL	25.7MG/100ML; 3.5GM/100ML; 5.08MG/100ML; 5 38MG/100ML; 448MG/100ML	N018379 008	Jun 24, 1988
-----------------	---	-------------	--------------

DELFLX-LM W/ DEXTROSE 4.25% IN PLASTIC CONTAINER

FRESENIUS MEDCL	25.7MG/100ML; 4.25GM/100ML; 5.08MG/100ML; 538MG/100ML; 448MG/100ML	N018379 006	Jul 07, 1982
-----------------	---	-------------	--------------

DIALYTE LM/ DEXTROSE 1.5% IN PLASTIC CONTAINER

B BRAUN	26MG/100ML; 1.5GM/100ML; 5MG/100ML; 530MG/ 100ML; 450MG/100ML	N018460 007	Jan 29, 1986
---------	--	-------------	--------------

	26MG/100ML; 1.5GM/100ML; 15MG/100ML; 560MG /100ML; 390MG/100ML	N018460 002	
--	---	-------------	--

DIALYTE LM/ DEXTROSE 2.5% IN PLASTIC CONTAINER

B BRAUN	26MG/100ML; 2.5GM/100ML; 5MG/100ML; 530MG/ 100ML; 450MG/100ML	N018460 005	Nov 02, 1983
---------	--	-------------	--------------

	26MG/100ML; 5GM/100ML; 5MG/100ML; 530MG/10 0ML; 450MG/100ML	N018460 008	Jan 29, 1986
--	--	-------------	--------------

DIALYTE LM/ DEXTROSE 4.25% IN PLASTIC CONTAINER

B BRAUN	26MG/100ML; 4.25GM/100ML; 5MG/100ML; 530MG /100ML; 450MG/100ML	N018460 009	Jan 29, 1986
---------	---	-------------	--------------

	26MG/100ML; 4.25GM/100ML; 15MG/100ML; 560M G/100ML; 390MG/100ML	N018460 004	
--	--	-------------	--

DIANEAL 137 W/ DEXTROSE 1.5% IN PLASTIC CONTAINER

BAXTER HLTHCARE	25.7MG/100ML; 1.5GM/100ML; 15.2MG/100ML; 5 67MG/100ML; 392MG/100ML	N017512 001	
-----------------	---	-------------	--

DIANEAL 137 W/ DEXTROSE 2.5% IN PLASTIC CONTAINER

BAXTER HLTHCARE	25.7MG/100ML; 2.5GM/100ML; 15.2MG/100ML; 5 67MG/100ML; 392MG/100ML	N017512 003	
-----------------	---	-------------	--

DIANEAL 137 W/ DEXTROSE 4.25% IN PLASTIC CONTAINER

BAXTER HLTHCARE	25.7MG/100ML; 4.25GM/100ML; 15.2MG/100ML; 567MG/100ML; 392MG/100ML	N017512 002	
-----------------	---	-------------	--

DIANEAL LOW CALCIUM W/ DEXTROSE 3.5% IN PLASTIC CONTAINER

+ BAXTER HLTHCARE	18.3MG/100ML; 3.5GM/100ML; 5.08MG/100ML; 5 38MG/100ML; 448MG/100ML	N020183 003	Dec 04, 1992
-------------------	---	-------------	--------------

DIANEAL PD-1 W/ DEXTROSE 1.5% IN PLASTIC CONTAINER

BAXTER HLTHCARE	25.7MG/100ML; 1.5GM/100ML; 15.2MG/100ML; 5 67MG/100ML; 392MG/100ML	N017512 007	Jul 09, 1984
-----------------	---	-------------	--------------

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CALCIUM CHLORIDE; DEXTROSE; MAGNESIUM CHLORIDE; SODIUM CHLORIDE; SODIUM LACTATE

SOLUTION; INTRAPERITONEAL

DIANEAL PD-1 W/ DEXTROSE 2.5% IN PLASTIC CONTAINER					
BAXTER HLTHCARE	25.7MG/100ML; 2.5GM/100ML; 15.2MG/100ML; 567MG/100ML; 392MG/100ML	N017512 008	Jul 09, 1984		
DIANEAL PD-1 W/ DEXTROSE 3.5% IN PLASTIC CONTAINER					
BAXTER HLTHCARE	25.7MG/100ML; 3.5GM/100ML; 15.2MG/100ML; 567MG/100ML; 392MG/100ML	N017512 010	Nov 18, 1985		
DIANEAL PD-1 W/ DEXTROSE 4.25% IN PLASTIC CONTAINER					
BAXTER HLTHCARE	25.7MG/100ML; 4.25GM/100ML; 15.2MG/100ML; 567MG/100ML; 392MG/100ML	N017512 009	Jul 09, 1984		
DIANEAL PD-2 W/ DEXTROSE 3.5% IN PLASTIC CONTAINER					
+ BAXTER HLTHCARE	25.7MG/100ML; 3.5GM/100ML; 5.08MG/100ML; 538MG/100ML; 448MG/100ML	N017512 011	Nov 18, 1985		
INPERSOL-LC/LM W/ DEXTROSE 1.5% IN PLASTIC CONTAINER					
FRESENIUS	18.4MG/100ML; 1.5GM/100ML; 5.08MG/100ML; 538MG/100ML; 448MG/100ML	A020374 001	Jun 13, 1994		
INPERSOL-LC/LM W/ DEXTROSE 2.5% IN PLASTIC CONTAINER					
FRESENIUS	18.4MG/100ML; 2.5GM/100ML; 5.08MG/100ML; 538MG/100ML; 448MG/100ML	A020374 002	Jun 13, 1994		
INPERSOL-IC/LM W/ DEXTROSE 3.5% IN PLASTIC CONTAINER					
FRESENIUS	18.4MG/100ML; 3.5GM/100ML; 5.08MG/100ML; 538MG/100ML; 448MG/100ML	A020374 003	Jun 13, 1994		
INPERSOL-LC/LM W/ DEXTROSE 4.25% IN PLASTIC CONTAINER					
FRESENIUS	18.4MG/100ML; 4.25GM/100ML; 5.08MG/100ML; 538MG/100ML; 448MG/100ML	A020374 004	Jun 13, 1994		

CALCIUM CHLORIDE; DEXTROSE; POTASSIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE

INJECTABLE; INJECTION

DEXTROSE 5% IN ACETATED RINGER'S IN PLASTIC CONTAINER					
B BRAUN	20MG/100ML; 5GM/100ML; 30MG/100ML; 380MG/100ML; 60MG/100ML	N018258 001			

CALCIUM CHLORIDE; DEXTROSE; POTASSIUM CHLORIDE; SODIUM CHLORIDE

INJECTABLE; INJECTION

DEXTROSE 5% AND RINGER'S IN PLASTIC CONTAINER					
HOSPIRA	33MG/100ML; 5GM/100ML; 30MG/100ML; 860MG/100ML	N018254 001			
DEXTROSE 5% IN RINGER'S IN PLASTIC CONTAINER					
B BRAUN	33MG/100ML; 5GM/100ML; 30MG/100ML; 860MG/100ML	N018256 001			
	33MG/100ML; 5GM/100ML; 30MG/100ML; 860MG/100ML	N020000 001	Apr 17, 1992		
BAXTER HLTHCARE	33MG/100ML; 5GM/100ML; 30MG/100ML; 860MG/100ML	N016695 001			

CALCIUM CHLORIDE; DEXTROSE; POTASSIUM CHLORIDE; SODIUM CHLORIDE; SODIUM LACTATE

INJECTABLE; INJECTION

DEXTROSE 4% IN MODIFIED LACTATED RINGER'S IN PLASTIC CONTAINER					
B BRAUN	4MG/100ML; 4GM/100ML; 6MG/100ML; 120MG/100ML; 62MG/100ML	N019634 002	Feb 24, 1988		
DEXTROSE 5% AND LACTATED RINGER'S					
FRESENIUS KABI USA	20MG/100ML; 5GM/100ML; 30MG/100ML; 600MG/100ML; 310MG/100ML	A210332 001	Mar 28, 2022		
DEXTROSE 5% IN LACTATED RINGER'S IN PLASTIC CONTAINER					
B BRAUN	20MG/100ML; 5GM/100ML; 30MG/100ML; 600MG/100ML; 310MG/100ML	N017510 001			
MILES	20MG/100ML; 5GM/100ML; 30MG/100ML; 600MG/100ML; 310MG/100ML	N018499 001			
POTASSIUM CHLORIDE 10MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER					
+ ICU MEDICAL INC	20MG/100ML; 5GM/100ML; 104MG/100ML; 600MG/100ML; 310MG/100ML	N019685 005	Oct 17, 1988		
+ ICU MEDICAL INC	20MG/100ML; 5GM/100ML; 179MG/100ML; 600MG/100ML; 310MG/100ML	N019685 006	Oct 17, 1988		
POTASSIUM CHLORIDE 15MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER					
+ ICU MEDICAL INC	20MG/100ML; 5GM/100ML; 254MG/100ML; 600MG/100ML; 310MG/100ML	N019685 007	Oct 17, 1988		
POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER					
FRESENIUS KABI USA	20MG/100ML; 5GM/100ML; 179MG/100ML; 600MG/100ML; 310MG/100ML	A211428 001	Nov 09, 2021		
+ ICU MEDICAL INC	20MG/100ML; 5GM/100ML; 328MG/100ML; 600MG/100ML; 310MG/100ML	N019685 008	Oct 17, 1988		
POTASSIUM CHLORIDE 30MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER					
+ ICU MEDICAL INC	20MG/100ML; 5GM/100ML; 254MG/100ML; 600MG/100ML; 310MG/100ML	N019685 003	Oct 17, 1988		

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CALCIUM CHLORIDE; DEXTROSE; POTASSIUM CHLORIDE; SODIUM CHLORIDE; SODIUM LACTATE

INJECTABLE; INJECTION

POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER			
+ ICU MEDICAL INC	20MG/100ML; 5GM/100ML; 328MG/100ML; 600MG/100ML; 310MG/100ML	N019685 004	Oct 17, 1988
POTASSIUM CHLORIDE 5MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER			
+ ICU MEDICAL INC	20MG/100ML; 5GM/100ML; 104MG/100ML; 600MG/100ML; 310MG/100ML	N019685 001	Oct 17, 1988

CALCIUM CHLORIDE; DEXTROSE; SODIUM CHLORIDE; SODIUM LACTATE

SOLUTION; INTRAPERITONEAL

INPERSOL-ZM W/ DEXTROSE 1.5% IN PLASTIC CONTAINER			
FRESENIUS MEDCL	25.7MG/100ML; 1.5GM/100ML; 538MG/100ML; 448MG/100ML	N019395 001	Mar 26, 1986
INPERSOL-ZM W/ DEXTROSE 2.5% IN PLASTIC CONTAINER			
FRESENIUS MEDCL	25.7MG/100ML; 2.5GM/100ML; 538MG/100ML; 448MG/100ML	N019395 002	Mar 26, 1986
INPERSOL-ZM W/ DEXTROSE 4.25% IN PLASTIC CONTAINER			
FRESENIUS MEDCL	25.7MG/100ML; 4.25GM/100ML; 538MG/100ML; 448MG/100ML	N019395 003	Mar 26, 1986

CALCIUM CHLORIDE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE

INJECTABLE; INJECTION

TPN ELECTROLYTES IN PLASTIC CONTAINER			
ABBOTT	16.5MG/ML; 25.4MG/ML; 74.6MG/ML; 121MG/ML; 16.1MG/ML	N019399 001	Jun 16, 1986

CALCIUM CHLORIDE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE; SODIUM CITRATE

INJECTABLE; INJECTION

ISOLYTE E IN PLASTIC CONTAINER			
B BRAUN	35MG/100ML; 30MG/100ML; 74MG/100ML; 640MG/100ML; 500MG/100ML; 74MG/100ML	N018899 001	Oct 31, 1983
	35MG/100ML; 30MG/100ML; 74MG/100ML; 640MG/100ML; 500MG/100ML; 74MG/100ML	N019718 001	Sep 29, 1989
SOLUTION; IRRIGATION			
BALANCED SALT			
EPIC PHARMA LLC	0.48MG/ML; 0.3MG/ML; 0.75MG/ML; 3.9MG/ML; 6.4MG/ML; 1.7MG/ML	A075503 001	Sep 27, 2006

CALCIUM CHLORIDE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE; SODIUM LACTATE

INJECTABLE; INJECTION

PLASMA-LYTE R IN PLASTIC CONTAINER			
BAXTER HLTHCARE	36.8MG/100ML; 30.5MG/100ML; 74.6MG/100ML; 640MG/100ML; 496MG/100ML; 89.6MG/100ML	N017438 001	

CALCIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE

INJECTABLE; INJECTION

ACETATED RINGER'S IN PLASTIC CONTAINER			
B BRAUN	20MG/100ML; 30MG/100ML; 380MG/100ML; 600MG/100ML	N018725 001	Nov 29, 1982

CALCIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM CHLORIDE

INJECTABLE; INJECTION

RINGER'S IN PLASTIC CONTAINER			
B BRAUN	33MG/100ML; 30MG/100ML; 860MG/100ML	N018721 001	Nov 09, 1982

SOLUTION; IRRIGATION

RINGER'S IN PLASTIC CONTAINER			
ABBOTT	33MG/100ML; 30MG/100ML; 860MG/100ML	N018462 001	

CALCIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM CHLORIDE; SODIUM LACTATE

INJECTABLE; INJECTION

LACTATED RINGER'S IN PLASTIC CONTAINER			
ABBOTT	20MG/100ML; 30MG/100ML; 600MG/100ML; 310MG/100ML	N019485 001	Oct 24, 1985
B BRAUN	20MG/100ML; 30MG/100ML; 600MG/100ML; 310MG/100ML	N018023 001	
MILES	20MG/100ML; 30MG/100ML; 600MG/100ML; 310MG/100ML	N018417 001	

SOLUTION; IRRIGATION

LACTATED RINGER'S IN PLASTIC CONTAINER			
BAXTER HLTHCARE	20MG/100ML; 30MG/100ML; 600MG/100ML; 310MG/100ML	N019933 001	Aug 29, 1989

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CALCIUM GLUCEPTATE

INJECTABLE; INJECTION

CALCIUM GLUCEPTATE

+ ABBOTT

EQ 90MG CALCIUM/5ML

A080001 001

EQ 90MG CALCIUM/5ML

A083159 001

ABRAXIS PHARM

EQ 90MG CALCIUM/5ML

A089373 001

Apr 30, 1987

LILLY

EQ 90MG CALCIUM/5ML

N006470 001

CALCIUM METRIZOATE; MEGLUMINE METRIZOATE; METRIZOATE MAGNESIUM; METRIZOATE SODIUM

INJECTABLE; INJECTION

ISOPAQUE 440

GE HEALTHCARE

0.78MG/ML; 75.9MG/ML; 0.15MG/ML; 16.6MG/ML

N016847 001

CALCIUM; MEGLUMINE; METRIZOIC ACID

INJECTABLE; INJECTION

ISOPAQUE 280

GE HEALTHCARE

0.35MG/ML; 140.1MG/ML; 461.8MG/ML

N017506 001

CANDESARTAN CILEXETIL

TABLET; ORAL

CANDESARTAN CILEXETIL

APOTEX

4MG

A202079 001 Jan 10, 2014

8MG

A202079 002 Jan 10, 2014

16MG

A202079 003 Jan 10, 2014

32MG

A202079 004 Jan 10, 2014

ZYDUS LIFESCIENCES

4MG

A091390 001 Aug 23, 2017

8MG

A091390 002 Aug 23, 2017

16MG

A091390 003 Aug 23, 2017

32MG

A091390 004 Aug 23, 2017

CANDESARTAN CILEXETIL; HYDROCHLOROTHIAZIDE

TABLET; ORAL

CANDESARTAN CILEXETIL AND HYDROCHLOROTHIAZIDE

APOTEX INC

16MG; 12.5MG

A202884 001 Dec 04, 2012

32MG; 12.5MG

A202884 002 Dec 04, 2012

32MG; 25MG

A202884 003 Jun 03, 2013

CANDICIDIN

OINTMENT; VAGINAL

VANOBIID

SANOFI AVENTIS US

0.6MG/GM

A061596 001

TABLET; VAGINAL

VANOBIID

SANOFI AVENTIS US

3MG

A061613 001

CAPECITABINE

TABLET; ORAL

CAPECITABINE

AMNEAL PHARMS

150MG

A204741 001 Feb 28, 2017

500MG

A204741 002 Feb 28, 2017

HIKMA

150MG

A200483 001 Jul 14, 2016

500MG

A200483 002 Jul 14, 2016

SUN PHARM

150MG

A204668 001 Jun 21, 2019

500MG

A204668 002 Jun 21, 2019

CAPREOMYCIN SULFATE

INJECTABLE; INJECTION

CAPASTAT SULFATE

+ EPIC PHARMA LLC

EQ 1GM BASE/VIAL **

N050095 001

CAPREOMYCIN SULFATE

HISUN PHARM HANGZHOU

EQ 1GM BASE/VIAL

A204796 001 Oct 18, 2018

NORVIUM BIOSCIENCE

EQ 1GM BASE/VIAL

A202634 001 Nov 27, 2017

CAPTOPRIL

TABLET; ORAL

CAPOTEN

+ STRIDES PHARMA

12.5MG **

N018343 005 Jan 17, 1985

+

25MG **

N018343 002

+

37.5MG **

N018343 006 Sep 17, 1986

+

50MG **

N018343 001

+

75MG **

N018343 007 Jun 13, 1995

+

100MG **

N018343 003

+

150MG **

N018343 004 Jun 13, 1995

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CAPTOPRIL

TABLET; ORAL

CAPTOPRIL

APOTHECON	12.5MG	A074472 001	Mar 31, 1995
	25MG	A074472 002	Mar 31, 1995
	50MG	A074472 003	Mar 31, 1995
	100MG	A074472 004	Mar 31, 1995
AUROBINDO PHARMA USA	12.5MG	A074434 001	Feb 13, 1996
	25MG	A074434 002	Feb 13, 1996
	50MG	A074434 003	Feb 13, 1996
	100MG	A074434 004	Feb 13, 1996
CHARTWELL RX	12.5MG	A074363 001	Nov 09, 1995
	12.5MG	A074519 001	Feb 13, 1996
	25MG	A074363 002	Nov 09, 1995
	25MG	A074519 002	Feb 13, 1996
	50MG	A074363 003	Nov 09, 1995
	50MG	A074519 003	Feb 13, 1996
	100MG	A074363 004	Nov 09, 1995
	100MG	A074519 004	Feb 13, 1996
COSETTE	12.5MG	A074433 001	Feb 13, 1996
	12.5MG	A074462 001	Feb 13, 1996
	12.5MG	A074483 001	Feb 13, 1996
	25MG	A074433 002	Feb 13, 1996
	25MG	A074462 002	Feb 13, 1996
	25MG	A074483 002	Feb 13, 1996
	50MG	A074433 003	Feb 13, 1996
	50MG	A074462 003	Feb 13, 1996
	50MG	A074483 003	Feb 13, 1996
	100MG	A074433 004	Feb 13, 1996
	100MG	A074462 004	Feb 13, 1996
	100MG	A074483 004	Feb 13, 1996
DAVA PHARMS INC	12.5MG	A074423 001	Feb 13, 1996
	25MG	A074423 002	Feb 13, 1996
	50MG	A074423 003	Feb 13, 1996
	100MG	A074423 004	Feb 13, 1996
EGIS PHARMS	12.5MG	A074748 004	May 29, 1997
	25MG	A074748 002	May 29, 1997
	50MG	A074748 001	May 29, 1997
	100MG	A074748 003	May 29, 1997
G AND W LABS INC	12.5MG	A074590 004	Aug 30, 1996
	25MG	A074590 002	Aug 30, 1996
	50MG	A074590 001	Aug 30, 1996
	100MG	A074590 003	Aug 30, 1996
OXFORD PHARMS	12.5MG	A074418 001	Feb 13, 1996
	25MG	A074418 002	Feb 13, 1996
	50MG	A074418 003	Feb 13, 1996
	100MG	A074418 004	Feb 13, 1996
PUREPAC PHARM	12.5MG	A074640 001	Mar 31, 1997
	25MG	A074640 002	Mar 31, 1997
	50MG	A074640 003	Mar 31, 1997
	100MG	A074640 004	Mar 31, 1997
SANDOZ	12.5MG	A074481 001	Feb 13, 1996
	25MG	A074481 002	Feb 13, 1996
	50MG	A074481 003	Feb 13, 1996
	100MG	A074481 004	Feb 13, 1996
SETON PHARMS	12.5MG	A212223 001	Oct 30, 2019
	25MG	A212223 002	Oct 30, 2019
	50MG	A212223 003	Oct 30, 2019
	100MG	A212223 004	Oct 30, 2019
STRIDES PHARMA	12.5MG	A074493 001	Feb 13, 1996
	25MG	A074493 002	Feb 13, 1996
	50MG	A074493 003	Feb 13, 1996
	100MG	A074493 004	Feb 13, 1996
TEVA	12.5MG	A074322 001	Feb 13, 1996
	25MG	A074322 002	Feb 13, 1996
	50MG	A074322 003	Feb 13, 1996
	100MG	A074322 004	Feb 13, 1996
WATSON LABS	12.5MG	A074386 001	May 23, 1996
	12.5MG	A074451 001	Feb 13, 1996

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CAPTOPRILTABLET; ORAL
CAPTOPRIL

12.5MG	A074576	001	Apr 23, 1996
25MG	A074386	002	May 23, 1996
25MG	A074451	002	Feb 13, 1996
25MG	A074576	002	Apr 23, 1996
50MG	A074386	003	May 23, 1996
50MG	A074451	003	Feb 13, 1996
50MG	A074576	003	Apr 23, 1996
100MG	A074386	004	May 23, 1996
100MG	A074451	004	Feb 13, 1996
100MG	A074576	004	Apr 23, 1996

CAPTOPRIL; HYDROCHLOROTHIAZIDE

TABLET; ORAL

CAPOZIDE 25/15

+ APOTHECON

25MG;15MG **

N018709 001 Oct 12, 1984

CAPOZIDE 25/25

+ APOTHECON

25MG;25MG **

N018709 002 Oct 12, 1984

CAPOZIDE 50/15

+ APOTHECON

50MG;15MG **

N018709 004 Oct 12, 1984

CAPOZIDE 50/25

+ APOTHECON

50MG;25MG **

N018709 003 Oct 12, 1984

CAPTOPRIL AND HYDROCHLOROTHIAZIDE

COSETTE

25MG;15MG

A074827 001 Dec 29, 1997

25MG;25MG

A074827 002 Dec 29, 1997

50MG;15MG

A074827 004 Dec 29, 1997

50MG;25MG

A074827 003 Dec 29, 1997

IVAX SUB TEVA PHARMS

25MG;15MG

A075055 001 Jun 18, 1998

25MG;25MG

A075055 002 Jun 18, 1998

50MG;15MG

A075055 004 Jun 18, 1998

50MG;25MG

A075055 003 Jun 18, 1998

STRIDES PHARMA

25MG;15MG

A074788 001 Dec 29, 1997

25MG;25MG

A074788 002 Dec 29, 1997

50MG;15MG

A074788 004 Dec 29, 1997

50MG;25MG

A074788 003 Dec 29, 1997

WATSON LABS

50MG;25MG

A074832 001 Dec 29, 1997

CARBACHOL

SOLUTION; INTRAOCULAR

CARBACHOL

PHARMAFAIR

0.01%

A070292 001 May 21, 1986

CARBASTAT

NOVARTIS

0.01%

A073677 001 Apr 28, 1995

CARBAMAZEPINE

SOLUTION; INTRAVENOUS

CARNEXIV

+ LUNDBECK PHARMS LLC

200MG/20ML (10MG/ML)

N206030 001 Oct 07, 2016

SUSPENSION; ORAL

CARBAMAZEPINE

TARO

100MG/5ML

A075875 001 Dec 21, 2000

TABLET; ORAL

CARBAMAZEPINE

ACTAVIS ELIZABETH

200MG

A071696 001 Nov 09, 1987

INWOOD LABS

200MG

A070231 001 Aug 14, 1986

PLIVA

200MG

A071479 001 Jul 24, 1987

RK PHARMA

200MG

A214328 001 Aug 16, 2021

TORRENT PHARMS

100MG

A077272 001 Dec 07, 2005

300MG

A077272 003 Dec 07, 2005

400MG

A077272 004 Dec 07, 2005

USL PHARMA

200MG

A070300 001 May 15, 1986

WARNER CHILCOTT

200MG

A070429 001 Jan 02, 1987

TERIL

TARO

200MG

A076525 001 Sep 26, 2003

TABLET, CHEWABLE; ORAL

CARBAMAZEPINE

JUBILANT CADISTA

100MG

A071940 001 Feb 01, 1988

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CARBAMAZEPINE

TABLET, CHEWABLE;ORAL

TEGRETOL

+ NOVARTIS

100MG **

N018281 001

TABLET, EXTENDED RELEASE;ORAL

CARBAMAZEPINE

AJANTA PHARMA LTD

100MG

A216193 001 Mar 24, 2023

200MG

A216193 002 Mar 24, 2023

400MG

A216193 003 Mar 24, 2023

AMNEAL PHARMS

100MG

A212704 001 Sep 22, 2023

200MG

A212704 002 Sep 22, 2023

400MG

A212704 003 Sep 22, 2023

APOTEX

100MG

A213159 001 Mar 08, 2024

200MG

A213159 002 Mar 08, 2024

400MG

A213159 003 Mar 08, 2024

CARBENICILLIN DISODIUM

INJECTABLE; INJECTION

GEOPEN

ROERIG

EQ 1GM BASE/VIAL

N050306 001

EQ 2GM BASE/VIAL

N050306 004

EQ 5GM BASE/VIAL

N050306 002

EQ 10GM BASE/VIAL

N050306 006

EQ 30GM BASE/VIAL

N050306 007

PYOPEN

GLAXOSMITHKLINE

EQ 1GM BASE/VIAL

N050298 001

EQ 2GM BASE/VIAL

N050298 002

EQ 5GM BASE/VIAL

N050298 003

EQ 10GM BASE/VIAL

N050298 006

EQ 20GM BASE/VIAL

N050298 007

CARBENICILLIN INDANYL SODIUM

TABLET;ORAL

GEOCILLIN

PFIZER

EQ 382MG BASE

N050435 001

CARBIDOPA

TABLET;ORAL

CARBIDOPA

ANI PHARMS

25MG

A203261 001 Mar 10, 2014

CARBIDOPA; ENTACAPONE; LEVODOPA

TABLET;ORAL

CARBIDOPA, LEVODOPA AND ENTACAPONE

NORVIUM BIOSCIENCE

12.5MG;200MG;50MG

A203424 001 Aug 13, 2020

18.75MG;200MG;75MG

A203424 002 Aug 13, 2020

25MG;200MG;100MG

A203424 003 Aug 13, 2020

31.25MG;200MG;125MG

A203424 004 Aug 13, 2020

37.5MG;200MG;150MG

A203424 005 Aug 13, 2020

50MG;200MG;200MG

A203424 006 Aug 13, 2020

WOCKHARDT LTD

12.5MG;200MG;50MG

A090786 001 Nov 20, 2012

18.75MG;200MG;75MG

A090833 001 Nov 20, 2012

25MG;200MG;100MG

A090833 002 Nov 20, 2012

31.25MG;200MG;125MG

A090833 003 Nov 20, 2012

37.5MG;200MG;150MG

A090833 004 Nov 20, 2012

50MG;200MG;200MG

A090833 005 Nov 20, 2012

CARBIDOPA; LEVODOPA

TABLET;ORAL

CARBIDOPA AND LEVODOPA

ANI PHARMS

10MG;100MG

A073587 002 Jun 29, 1995

25MG;100MG

A073587 001 Jun 29, 1995

25MG;250MG

A073587 003 Jun 29, 1995

SCS

10MG;100MG

A074080 001 Mar 25, 1994

25MG;100MG

A074080 002 Mar 25, 1994

25MG;250MG

A074080 003 Mar 25, 1994

WATSON LABS

10MG;100MG

A073381 001 Sep 28, 1993

25MG;100MG

A073382 001 Sep 28, 1993

25MG;250MG

A073383 001 Sep 28, 1993

ZYDUS PHARMS

10MG;100MG

A215999 001 Apr 04, 2023

25MG;100MG

A215999 002 Apr 04, 2023

25MG;250MG

A215999 003 Apr 04, 2023

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CARBIDOPA; LEVODOPA

TABLET, EXTENDED RELEASE;ORAL

CARBIDOPA AND LEVODOPA

KV PHARM

50MG;200MG

A076663 001 Jun 24, 2004

SINEMET CR

+ ORGANON LLC

25MG;100MG **

N019856 002 Dec 24, 1992

+

50MG;200MG **

N019856 001 May 30, 1991

TABLET, FOR SUSPENSION;ORAL

CARBILEV

RANBAXY

10MG;100MG

A076643 001 Jun 10, 2005

25MG;100MG

A076643 002 Jun 10, 2005

25MG;250MG

A076643 003 Jun 10, 2005

TABLET, ORALLY DISINTEGRATING;ORAL

CARBIDOPA AND LEVODOPA

IMPAX LABS

10MG;100MG

A090631 001 Jun 08, 2010

25MG;100MG

A090631 002 Jun 08, 2010

25MG;250MG

A090631 003 Jun 08, 2010

RISING

10MG;100MG

A078893 001 Sep 18, 2008

25MG;100MG

A078893 002 Sep 18, 2008

25MG;250MG

A078893 003 Sep 18, 2008

PARCOPA

UCB INC

10MG;100MG **

A076699 001 Aug 27, 2004

25MG;100MG **

A076699 002 Aug 27, 2004

25MG;250MG **

A076699 003 Aug 27, 2004

CARBINOXAMINE MALEATE

ELIXIR;ORAL

CLISTIN

+ MCNEIL

4MG/5ML **

N008955 001

SOLUTION;ORAL

CARBINOXAMINE MALEATE

CYPRESS PHARM

4MG/5ML

A090418 001 May 04, 2010

ENDO OPERATIONS

4MG/5ML

A040814 001 Feb 26, 2008

TABLET;ORAL

CARBINOXAMINE MALEATE

CYPRESS PHARM

4MG

A090417 001 Aug 23, 2010

INVAGEN PHARMS

4MG

A090435 001 Apr 15, 2010

STRIDES PHARMA

4MG

A040639 002 May 30, 2008

CLISTIN

+ ORTHO MCNEIL PHARM

4MG **

N008915 001

CARBOPLATIN

INJECTABLE; INJECTION

CARBOPLATIN

CIPLA LTD

50MG/VIAL

A077383 001 Jan 27, 2006

150MG/VIAL

A077383 002 Jan 27, 2006

450MG/VIAL

A077383 003 Jan 27, 2006

FRESENIUS KABI USA

50MG/VIAL

A076235 001 Oct 14, 2004

150MG/VIAL

A076235 002 Oct 14, 2004

450MG/VIAL

A076235 003 Oct 14, 2004

HIKMA

50MG/VIAL

A076099 001 Oct 14, 2004

150MG/VIAL

A076099 002 Oct 14, 2004

450MG/VIAL

A076099 003 Oct 14, 2004

HOSPIRA

50MG/VIAL

A076473 001 Oct 27, 2004

150MG/VIAL

A076473 002 Oct 27, 2004

450MG/VIAL

A076473 003 Oct 27, 2004

NATCO PHARMA USA

50MG/VIAL

A091510 001 May 29, 2012

150MG/VIAL

A091510 002 May 29, 2012

450MG/VIAL

A091510 003 May 29, 2012

PLIVA

50MG/VIAL

A076602 001 Nov 16, 2004

150MG/VIAL

A076602 002 Nov 16, 2004

450MG/VIAL

A076602 003 Nov 16, 2004

SANDOZ

50MG/VIAL

A076959 001 Mar 18, 2005

150MG/VIAL

A076959 002 Mar 18, 2005

450MG/VIAL

A076959 003 Mar 18, 2005

WATSON LABS TEVA

50MG/VIAL

A076162 001 Oct 14, 2004

150MG/VIAL

A076162 002 Oct 14, 2004

450MG/VIAL

A076162 003 Oct 14, 2004

PARAPLATIN

+ CORDEN PHARMA

50MG/VIAL **

N019880 001 Mar 03, 1989

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CARBOPLATIN

INJECTABLE; INJECTION

PARAPLATIN

+

150MG/VIAL **

N019880 002 Mar 03, 1989

+

450MG/VIAL **

N019880 003 Mar 03, 1989

INJECTABLE; INTRAVENOUS

CARBOPLATIN

ACTAVIS TOTOWA

50MG/5ML (10MG/ML)

A078732 001 Feb 06, 2012

150MG/15ML (10MG/ML)

A078732 002 Feb 06, 2012

450MG/45ML (10MG/ML)

A078732 003 Feb 06, 2012

600MG/60ML (10MG/ML)

A078732 004 Feb 06, 2012

FRESENIUS KABI USA

50MG/5ML (10MG/ML)

A077432 001 Sep 29, 2006

150MG/15ML (10MG/ML)

A077432 002 Sep 29, 2006

450MG/45ML (10MG/ML)

A077432 003 Sep 29, 2006

50MG/5ML (10MG/ML)

A077247 001 Oct 21, 2004

50MG/5ML (10MG/ML)

A077266 001 Feb 15, 2006

150MG/15ML (10MG/ML)

A077247 002 Oct 21, 2004

150MG/15ML (10MG/ML)

A077266 002 Feb 15, 2006

MEITHEAL

50MG/5ML (10MG/ML)

A077096 001 Jun 14, 2005

150MG/15ML (10MG/ML)

A077096 002 Jun 14, 2005

450MG/45ML (10MG/ML)

A077096 003 Jun 14, 2005

600MG/60ML (10MG/ML)

A077096 004 Jun 03, 2013

NORVIUM BIOSCIENCE

50MG/5ML (10MG/ML)

A077998 001 Apr 24, 2007

50MG/5ML (10MG/ML)

A091063 001 Nov 09, 2011

150MG/15ML (10MG/ML)

A077998 002 Apr 24, 2007

150MG/15ML (10MG/ML)

A091063 002 Nov 09, 2011

450MG/45ML (10MG/ML)

A077998 003 Apr 24, 2007

450MG/45ML (10MG/ML)

A091063 003 Nov 09, 2011

600MG/60ML (10MG/ML)

A091063 004 Nov 09, 2011

1GM/100ML (10MG/ML)

A091478 001 Nov 23, 2011

PHARMACHEMIE BV

50MG/5ML (10MG/ML)

A077679 001 Feb 25, 2009

150MG/15ML (10MG/ML)

A077679 002 Feb 25, 2009

450MG/45ML (10MG/ML)

A077679 003 Feb 25, 2009

PLIVA LACHEMA

50MG/5ML (10MG/ML)

A078631 001 Dec 02, 2008

150MG/15ML (10MG/ML)

A078631 002 Dec 02, 2008

450MG/45ML (10MG/ML)

A078631 003 Dec 02, 2008

600MG/60ML (10MG/ML)

A078631 004 Dec 02, 2008

TEVA PARENTERAL

50MG/5ML (10MG/ML)

A077389 001 Mar 30, 2007

150MG/15ML (10MG/ML)

A077389 002 Mar 30, 2007

450MG/45ML (10MG/ML)

A077389 003 Mar 30, 2007

TEVA PHARMS USA

50MG/5ML (10MG/ML)

A077139 001 Sep 21, 2005

150MG/15ML (10MG/ML)

A077139 002 Sep 21, 2005

450MG/45ML (10MG/ML)

A077139 003 Sep 21, 2005

600MG/60ML (10MG/ML)

A077139 004 Sep 21, 2005

PARAPLATIN

+ CORDENPHARMA

50MG/5ML (10MG/ML) **

N020452 001 Jul 14, 2003

+

150MG/15ML (10MG/ML) **

N020452 002 Jul 14, 2003

+

450MG/45ML (10MG/ML) **

N020452 003 Jul 14, 2003

+

600MG/60ML (10MG/ML) **

N020452 004 Jan 15, 2004

CARBOPROST TROMETHAMINE

INJECTABLE; INJECTION

CARBOPROST TROMETHAMINE

ANI PHARMS

EQ 0.25MG BASE/ML

A215901 001 Jan 25, 2024

CARFILZOMIB

POWDER; INTRAVENOUS

CARFILZOMIB

APOTEX

30MG/VIAL

A211185 001 Mar 20, 2020

60MG/VIAL

A209425 001 Mar 16, 2020

BRECKENRIDGE

10MG/VIAL

A209330 001 Jun 11, 2021

60MG/VIAL

A209330 002 Jun 11, 2021

CARIPRAZINE HYDROCHLORIDE

CAPSULE; ORAL

CARIPRAZINE HYDROCHLORIDE

SUN PHARM

EQ 1.5MG BASE

A213932 001 Sep 30, 2022

EQ 3MG BASE

A213932 002 Sep 30, 2022

EQ 4.5MG BASE

A213932 003 Sep 30, 2022

EQ 6MG BASE

A213932 004 Sep 30, 2022

ZYDUS

EQ 1.5MG BASE

A213984 001 Sep 09, 2022

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CARIPRAZINE HYDROCHLORIDE

CAPSULE; ORAL

CARIPRAZINE HYDROCHLORIDE

EQ 3MG BASE	A213984 002	Sep 09, 2022
EQ 4.5MG BASE	A213984 003	Sep 09, 2022
EQ 6MG BASE	A213984 004	Sep 09, 2022

CARISOPRODOL

CAPSULE; ORAL

SOMA

+ MYLAN SPECIALITY LP 250MG

N011792 003

TABLET; ORAL

CARISOPRODOL

ABLE	350MG	A040421 001	Jun 21, 2001
EPIC PHARMA LLC	350MG	A040397 001	Sep 21, 2000
HIKMA INTL PHARMS	350MG	A040124 001	Jan 24, 1996
PIONEER PHARMS	350MG	A089390 001	Oct 13, 1988
SANDOZ	350MG	A089566 001	Aug 30, 1988
STRIDES PHARMA	250MG	A205513 001	Nov 12, 2015
	350MG	A205513 002	Nov 12, 2015
SUN PHARM INDS LTD	350MG	A040755 001	Feb 27, 2007
SUN PHARM INDUSTRIES	350MG	A089346 001	Oct 17, 1991
WATSON LABS	350MG	A040152 001	Dec 03, 1996
	350MG	A085433 001	
WATSON LABS TEVA	350MG	A086179 001	

RELA

SCHERING	350MG	N012155 001	
----------	-------	-------------	--

CARMUSTINE

INJECTABLE; INJECTION

CARMUSTINE

ACCORD HLTHCARE	100MG/VIAL	A214117 001	Dec 27, 2022
INGENUS PHARMS LLC	100MG/VIAL	A211011 001	Jun 21, 2022
NORVIUM BIOSCIENCE	100MG/VIAL	A215368 001	Mar 03, 2023

POWDER; INTRAVENOUS

CARMUSTINE

+ ACCORD HLTHCARE	50MG/VIAL	N215000 001	May 16, 2022
+	300MG/VIAL	N215000 002	May 16, 2022

CARPENAZINE MALEATE

CONCENTRATE; ORAL

PROKETAZINE

WYETH AYERST	50MG/ML	N014173 001	
--------------	---------	-------------	--

TABLET; ORAL

PROKETAZINE

WYETH AYERST	12.5MG	N012768 001	
	25MG	N012768 002	
	50MG	N012768 004	

CARPROFEN

TABLET; ORAL

RIMADYL

ROCHE	100MG	N018550 002	Dec 31, 1987
	150MG	N018550 003	Dec 31, 1987

CARTEOLOL HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

CARTEOLOL HYDROCHLORIDE

APOTEX INC	1%	A076097 001	Feb 06, 2002
BAUSCH AND LOMB	1%	A075546 001	Jan 20, 2000

OCUPRESS

+ NOVARTIS	1% **	N019972 001	May 23, 1990
------------	-------	-------------	--------------

TABLET; ORAL

CARTROL

ABBVIE	2.5MG	N019204 001	Dec 28, 1988
	5MG	N019204 002	Dec 28, 1988
	10MG	N019204 003	Dec 28, 1988

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CARVEDILOL

TABLET; ORAL

CARVEDILOL

HIKMA	3.125MG	A077887 001	Sep 07, 2007
	6.25MG	A077887 002	Sep 07, 2007
	12.5MG	A077887 003	Sep 07, 2007
	25MG	A077887 004	Sep 07, 2007
MYLAN	3.125MG	A077316 001	Sep 05, 2007
PLIVA HRVATSKA DOO	3.125MG	A078240 001	Oct 30, 2007
	6.25MG	A078240 002	Oct 30, 2007
	12.5MG	A078240 003	Oct 30, 2007
	25MG	A078240 004	Oct 30, 2007
SUN PHARM INDS INC	3.125MG	A077346 004	Sep 05, 2007
	6.25MG	A077346 001	Sep 05, 2007
	12.5MG	A077346 002	Sep 05, 2007
	25MG	A077346 003	Sep 05, 2007
SUN PHARM INDS LTD	3.125MG	A076989 001	Sep 05, 2007
	6.25MG	A076989 002	Sep 05, 2007
	12.5MG	A076989 003	Sep 05, 2007
	25MG	A076989 004	Sep 05, 2007

CARVEDILOL PHOSPHATE

CAPSULE, EXTENDED RELEASE; ORAL

COREG CR

+ WAYLIS THERAP	10MG	N022012 001	Oct 20, 2006
+	20MG	N022012 002	Oct 20, 2006
+	40MG	N022012 003	Oct 20, 2006
+	80MG	N022012 004	Oct 20, 2006

CASPOFUNGIN ACETATE

POWDER; INTRAVENOUS

CANCIDAS

+ MERCK	50MG/VIAL	N021227 001	Jan 26, 2001
+	70MG/VIAL	N021227 002	Jan 26, 2001

CASPOFUNGIN ACETATE

CIPLA	50MG/VIAL	A209489 001	Jul 12, 2018
	70MG/VIAL	A209489 002	Jul 12, 2018
NORVIUM BIOSCIENCE	50MG/VIAL	A207650 001	Sep 29, 2017
	70MG/VIAL	A207650 002	Sep 29, 2017
XELLIA PHARMS APS	50MG/VIAL	A205923 001	Jul 02, 2018
	70MG/VIAL	A205923 002	Jul 02, 2018

CEFACLOR

CAPSULE; ORAL

CECLOR

+ LILLY	EQ 250MG BASE **	N050521 001	
+	EQ 500MG BASE **	N050521 002	

CEFACLOR

CEPH INTL	EQ 250MG BASE	A062205 001	
	EQ 500MG BASE	A062205 002	
CHARTWELL RX	EQ 250MG BASE	A064148 001	May 23, 1996
	EQ 500MG BASE	A064148 002	May 23, 1996
DAVA PHARMS INC	EQ 250MG BASE	A064107 001	Apr 27, 1995
	EQ 500MG BASE	A064107 002	Apr 27, 1995
HIKMA	EQ 250MG BASE	A065350 001	Apr 03, 2007
	EQ 500MG BASE	A065350 002	Apr 03, 2007
IVAX SUB TEVA PHARMS	EQ 250MG BASE	A064061 001	Apr 27, 1995
	EQ 500MG BASE	A064061 002	Apr 27, 1995
RANBAXY	EQ 250MG BASE	A064156 001	Aug 28, 1997
	EQ 500MG BASE	A064156 002	Aug 28, 1997
TEVA	EQ 250MG BASE	A064081 001	Sep 16, 1996
	EQ 250MG BASE	A064145 001	Jun 24, 1996
	EQ 500MG BASE	A064081 002	Sep 16, 1996
	EQ 500MG BASE	A064145 002	Jun 24, 1996

FOR SUSPENSION; ORAL

CECLOR

+ LILLY	EQ 125MG BASE/5ML **	N050522 001	
+	EQ 250MG BASE/5ML **	N050522 002	

CEFACLOR

DAVA PHARMS INC	EQ 125MG BASE/5ML	A064114 001	Apr 28, 1995
	EQ 187MG BASE/5ML	A064115 001	Apr 28, 1995

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CEFACTOR

FOR SUSPENSION;ORAL

CEFACTOR

	EQ 250MG BASE/5ML	A064116 001	Apr 28, 1995
	EQ 375MG BASE/5ML	A064110 001	Apr 28, 1995
FACTA FARMA	EQ 125MG BASE/5ML	A062206 001	
	EQ 187MG BASE/5ML	A062206 003	Apr 20, 1988
	EQ 250MG BASE/5ML	A062206 002	
	EQ 375MG BASE/5ML	A062206 004	Apr 20, 1988
IVAX SUB TEVA PHARMS	EQ 125MG BASE/5ML	A064087 001	Apr 28, 1995
	EQ 187MG BASE/5ML	A064086 001	Apr 28, 1995
	EQ 250MG BASE/5ML	A064085 001	Apr 28, 1995
	EQ 375MG BASE/5ML	A064070 001	Apr 28, 1995
RANBAXY	EQ 125MG BASE/5ML	A064166 001	Oct 02, 1997
	EQ 187MG BASE/5ML	A064165 001	Oct 02, 1997
	EQ 250MG BASE/5ML	A064164 001	Oct 02, 1997
	EQ 375MG BASE/5ML	A064155 001	Oct 02, 1997
WATSON LABS INC	EQ 125MG BASE/5ML	A064204 001	Feb 18, 1998
	EQ 187MG BASE/5ML	A064205 001	Feb 18, 1998
	EQ 250MG BASE/5ML	A064206 001	Feb 18, 1998
	EQ 375MG BASE/5ML	A064207 001	Feb 18, 1998

TABLET, CHEWABLE;ORAL

RANICLOR

RANBAXY LABS LTD	EQ 125MG BASE	A065092 001	Dec 22, 2003
	EQ 187MG BASE	A065092 002	Dec 22, 2003
	EQ 250MG BASE	A065092 003	Dec 22, 2003
	EQ 375MG BASE	A065092 004	Dec 22, 2003

TABLET, EXTENDED RELEASE;ORAL

CECLOR CD

LILLY	EQ 375MG BASE **	N050673 001	Jun 28, 1996
	EQ 500MG BASE **	N050673 002	Jun 28, 1996

CEFACTOR

WORLD GEN	EQ 500MG BASE	A065057 001	Jan 05, 2001
-----------	---------------	-------------	--------------

CEFADROXIL/CEFADROXIL HEMIHYDRATE

CAPSULE;ORAL

CEFADROXIL

CHARTWELL RX	EQ 500MG BASE	A065309 001	Sep 18, 2006
CSPC OUYI	EQ 500MG BASE	A205072 001	Jul 28, 2017
HIKMA	EQ 500MG BASE	A065311 001	Feb 07, 2006
IVAX SUB TEVA PHARMS	EQ 500MG BASE	A062766 001	Mar 03, 1987
PUREPAC PHARM	EQ 500MG BASE	A063017 001	Jan 05, 1989
RANBAXY LABS LTD	EQ 500MG BASE	A065015 001	Jun 22, 1999
SANDOZ	EQ 500MG BASE	A062291 001	
TEVA	EQ 500MG BASE	A062695 001	Feb 10, 1989

DURICEF

WARNER CHILCOTT	EQ 250MG BASE	N050512 002	
+	EQ 500MG BASE **	N050512 001	

ULTRACEF

BRISTOL	EQ 500MG BASE	A062378 001	Mar 16, 1982
---------	---------------	-------------	--------------

FOR SUSPENSION;ORAL

CEFADROXIL

ANI PHARMS	EQ 125MG BASE/5ML	A062698 001	Mar 01, 1989
	EQ 250MG BASE/5ML	A062698 002	Mar 01, 1989
	EQ 250MG BASE/5ML	A065278 001	Jan 20, 2006
	EQ 500MG BASE/5ML	A062698 003	Mar 01, 1989
	EQ 500MG BASE/5ML	A065278 002	Jan 20, 2006
APOTHECON	EQ 125MG BASE/5ML	A062334 001	
	EQ 250MG BASE/5ML	A062334 002	
	EQ 500MG BASE/5ML	A062334 003	
CHARTWELL RX	EQ 250MG BASE/5ML	A065307 002	Oct 16, 2006
	EQ 500MG BASE/5ML	A065307 003	Oct 16, 2006
HIKMA PHARMS	EQ 250MG BASE/5ML	A091036 001	Nov 28, 2012
	EQ 500MG BASE/5ML	A091036 002	Nov 28, 2012
SUN PHARM INDS LTD	EQ 125MG BASE/5ML	A065115 001	Mar 26, 2003
	EQ 250MG BASE/5ML	A065115 002	Mar 26, 2003
	EQ 500MG BASE/5ML	A065115 003	Mar 26, 2003

DURICEF

+	WARNER CHILCOTT	EQ 125MG BASE/5ML **	N050527 002
+		EQ 250MG BASE/5ML **	N050527 003

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CEFADROXIL/CEFADROXIL HEMIHYDRATE

FOR SUSPENSION;ORAL

DURICEF

+

EQ 500MG BASE/5ML **

N050527 001

ULTRACEF

BRISTOL

EQ 125MG BASE/5ML

A062376 001 Mar 16, 1982

EQ 250MG BASE/5ML

A062376 002 Mar 16, 1982

EQ 500MG BASE/5ML

A062376 003 Mar 16, 1982

TABLET;ORAL

CEFADROXIL

CHARTWELL RX

EQ 1GM BASE

A065301 001 Sep 18, 2006

HIKMA

EQ 1GM BASE

A065260 001 Mar 30, 2006

RANBAXY

EQ 1GM BASE

A065018 001 Apr 23, 1999

DURICEF

+

WARNER CHILCOTT

EQ 1GM BASE **

N050528 001

ULTRACEF

APOTHECON

EQ 1GM BASE

A062390 001 Jun 10, 1982

BRISTOL

EQ 1GM BASE

A062408 001 Aug 31, 1982

CEFAMANDOLE NAFATE

INJECTABLE;INJECTION

MANDOL

LILLY

EQ 500MG BASE/VIAL

N050504 001

EQ 1GM BASE/VIAL

A062560 001 Sep 10, 1985

EQ 1GM BASE/VIAL

N050504 002

EQ 2GM BASE/VIAL

A062560 002 Sep 10, 1985

EQ 2GM BASE/VIAL

N050504 003

EQ 10GM BASE/VIAL

N050504 004

CEFAZOLIN SODIUM

INJECTABLE;INJECTION

ANCEF

+

GLAXOSMITHKLINE

EQ 250MG BASE/VIAL **

N050461 001

+

EQ 500MG BASE/VIAL **

N050461 002

+

EQ 1GM BASE/VIAL **

N050461 003

+

EQ 5GM BASE/VIAL **

N050461 004

+

EQ 10GM BASE/VIAL **

N050461 005

ANCEF IN DEXTROSE 5% IN PLASTIC CONTAINER

+

BAXTER HLTHCARE

EQ 10MG BASE/ML

N050566 003 Jun 08, 1983

+

EQ 20MG BASE/ML

N050566 004 Jun 08, 1983

ANCEF IN PLASTIC CONTAINER

BAXTER HLTHCARE

EQ 10MG BASE/ML

A063002 001 Mar 28, 1991

EQ 20MG BASE/ML

A063002 002 Mar 28, 1991

ANCEF IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

+

BAXTER HLTHCARE

EQ 10MG BASE/ML

N050566 001 Jun 08, 1983

+

EQ 20MG BASE/ML

N050566 002 Jun 08, 1983

CEFAZOLIN AND DEXTROSE

B BRAUN

EQ 500MG BASE/VIAL

N050779 001 Jul 27, 2000

CEFAZOLIN SODIUM

ABRAXIS PHARM

EQ 500MG BASE/VIAL

A062688 002 Nov 17, 1986

EQ 1GM BASE/VIAL

A062688 003 Nov 17, 1986

EQ 10GM BASE/VIAL

A062688 004 Nov 17, 1986

EQ 20GM BASE/VIAL

A062688 005 Aug 03, 1987

AUROBINDO PHARMA

EQ 500MG BASE/VIAL

A065395 001 Aug 08, 2008

EQ 1GM BASE/VIAL

A065395 002 Aug 08, 2008

BEDFORD

EQ 250MG BASE/VIAL

A062894 001 Jul 21, 1988

EQ 500MG BASE/VIAL

A062894 002 Jul 21, 1988

EQ 1GM BASE/VIAL

A062894 003 Jul 21, 1988

EQ 5GM BASE/VIAL

A062894 004 Jul 21, 1988

EQ 10GM BASE/VIAL

A062894 005 Jul 21, 1988

CEPHAZONE PHARMA

EQ 500MG BASE/VIAL

A065280 001 Mar 18, 2009

EQ 1GM BASE/VIAL

A065280 002 Mar 18, 2009

EQ 10GM BASE/VIAL

A065295 001 Mar 18, 2009

EQ 20GM BASE/VIAL

A065296 001 Mar 18, 2009

DR REDDYS

EQ 250MG BASE/VIAL

A062988 001 Dec 29, 1989

EQ 500MG BASE/VIAL

A062988 002 Dec 29, 1989

EQ 1GM BASE/VIAL

A062988 003 Dec 29, 1989

EQ 5GM BASE/VIAL

A062989 001 Dec 29, 1989

EQ 10GM BASE/VIAL

A062989 002 Dec 29, 1989

EQ 20GM BASE/VIAL

A062989 003 Dec 29, 1989

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CEFAZOLIN SODIUM

INJECTABLE; INJECTION

CEFAZOLIN SODIUM

FACTA FARMA	EQ 500MG BASE/VIAL	A063214 001	Dec 27, 1991
	EQ 1GM BASE/VIAL	A063207 001	Dec 27, 1991
	EQ 10GM BASE/VIAL	A063209 001	Dec 27, 1991
	EQ 20GM BASE/VIAL	A063209 002	Apr 30, 1999
FRESENIUS KABI USA	EQ 500MG BASE/VIAL **	A064169 001	Aug 14, 1998
	EQ 1GM BASE/VIAL **	A064169 002	Aug 14, 1998
	EQ 10GM BASE/VIAL **	A064170 001	Mar 18, 1998
	EQ 20GM BASE/VIAL **	A064170 002	Mar 18, 1998
GLAXOSMITHKLINE	EQ 1GM BASE/VIAL	A064033 001	Oct 31, 1993
HIKMA	EQ 250MG BASE/VIAL	A062807 001	Jan 12, 1988
	EQ 500MG BASE/VIAL	A062807 002	Jan 12, 1988
	EQ 1GM BASE/VIAL	A062807 003	Jan 12, 1988
	EQ 5GM BASE/VIAL	A062807 004	Jan 12, 1988
	EQ 10GM BASE/VIAL	A062807 005	Jan 12, 1988
	EQ 20GM BASE/VIAL	A062807 006	Jan 12, 1988
HOSPIRA	EQ 1GM BASE/VIAL	A201654 001	Feb 03, 2016
HOSPIRA INC	EQ 500MG BASE/VIAL	A065226 001	Apr 21, 2005
	EQ 1GM BASE/VIAL	A065226 002	Apr 21, 2005
	EQ 1GM BASE/VIAL	A065244 001	Aug 12, 2005
	EQ 10GM BASE/VIAL	A065247 001	Aug 12, 2005
STERI PHARMA	EQ 500MG BASE/VIAL	A063216 001	Dec 27, 1991
	EQ 1GM BASE/VIAL	A063208 001	Dec 27, 1991
TEVA PHARMS	EQ 250MG BASE/VIAL	A063016 001	Mar 14, 1989
	EQ 500MG BASE/VIAL	A063016 002	Mar 14, 1989
	EQ 1GM BASE/VIAL	A063016 003	Mar 14, 1989
	EQ 5GM BASE/VIAL	A063018 001	Mar 05, 1990
	EQ 10GM BASE/VIAL	A063018 002	Mar 05, 1990
KEFZOL			
ACS DOBFAR	EQ 250MG BASE/VIAL	A061773 001	
	EQ 500MG BASE/VIAL	A061773 002	
	EQ 1GM BASE/VIAL	A061773 003	
	EQ 10GM BASE/VIAL	A061773 004	
	EQ 20GM BASE/VIAL	A061773 005	Sep 08, 1987
LILLY	EQ 500MG BASE/VIAL	A062557 001	Sep 10, 1985
	EQ 1GM BASE/VIAL	A062557 002	Sep 10, 1985

CEFDINIR

CAPSULE; ORAL

OMNICEF

+ ABBVIE

300MG **

N050739 001 Dec 04, 1997

FOR SUSPENSION; ORAL

OMNICEF

+ ABBVIE

125MG/5ML **

N050749 001 Dec 04, 1997

+

250MG/5ML **

N050749 002 Jul 29, 2004

CEFDITOREN PIVOXIL

TABLET; ORAL

SPECTRACEF

VANSEN PHARMA

200MG

N021222 001 Aug 29, 2001

400MG

N021222 002 Jul 21, 2008

CEFEPIME HYDROCHLORIDE

INJECTABLE; INJECTION

CEFEPIME HYDROCHLORIDE

ASTRAL

EQ 1GM BASE/VIAL

A212721 001 Jul 21, 2020

EQ 2GM BASE/VIAL

A212721 002 Jul 21, 2020

CHARTWELL RX

EQ 500MG BASE/VIAL

A090291 001 Dec 21, 2010

EQ 1GM BASE/VIAL

A090291 002 Dec 21, 2010

EQ 2GM BASE/VIAL

A090291 003 Dec 21, 2010

HOSPIRA INC

EQ 500MG BASE/VIAL

A065369 001 Jun 18, 2007

EQ 1GM BASE/VIAL

A065369 002 Jun 18, 2007

EQ 1GM BASE/VIAL

A202268 001 Jul 30, 2012

EQ 2GM BASE/VIAL

A065369 003 Jun 18, 2007

EQ 2GM BASE/VIAL

A202268 002 Jul 30, 2012

MAXIPIME

+ HOSPIRA INC

EQ 500MG BASE/VIAL **

N050679 001 Jan 18, 1996

+

EQ 1GM BASE/VIAL **

N050679 002 Jan 18, 1996

+

EQ 2GM BASE/VIAL **

N050679 003 Jan 18, 1996

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CEFEPIME HYDROCHLORIDE; ENMETAZOBACTAM

POWDER; INTRAVENOUS

EXBLIFEP

+ ALLECRA THERAPS EQ 2GM BASE/VIAL;0.5GM/VIAL N216165 001 Feb 22, 2024

CEFIXIME

FOR SUSPENSION; ORAL

CEFIXIME

CHARTWELL RX 100MG/5ML A206144 001 Nov 17, 2017

200MG/5ML A206144 002 Nov 17, 2017

SUPRAX

+ LEDERLE 100MG/5ML ** N050622 001 Apr 28, 1989

LUPIN PHARMS 100MG/5ML A065129 001 Feb 23, 2004

TABLET; ORAL

SUPRAX

+ LEDERLE 200MG ** N050621 001 Apr 28, 1989

+ 400MG ** N050621 002 Apr 28, 1989

LUPIN PHARMS 400MG A065130 001 Feb 12, 2004

CEFMENOXIME HYDROCHLORIDE

INJECTABLE; INJECTION

CEFMAX

TAP PHARM EQ 500MG BASE/VIAL N050571 001 Dec 30, 1987

EQ 1GM BASE/VIAL N050571 002 Dec 30, 1987

EQ 2GM BASE/VIAL N050571 003 Dec 30, 1987

CEFMETAZOLE SODIUM

INJECTABLE; INJECTION

ZEFAZONE

+ PHARMACIA AND UPJOHN EQ 1GM BASE/VIAL ** N050637 001 Dec 11, 1989

+ EQ 2GM BASE/VIAL ** N050637 002 Dec 11, 1989

ZEFAZONE IN PLASTIC CONTAINER

+ PHARMACIA AND UPJOHN EQ 20MG BASE/ML ** N050683 001 Dec 29, 1992

+ EQ 40MG BASE/ML ** N050683 002 Dec 29, 1992

CEFONICID SODIUM

INJECTABLE; INJECTION

MONOCID

GLAXOSMITHKLINE EQ 500MG BASE/VIAL N050579 001 May 23, 1984

EQ 1GM BASE/VIAL A063295 001 Jul 26, 1993

EQ 1GM BASE/VIAL N050579 002 May 23, 1984

EQ 2GM BASE/VIAL N050579 003 May 23, 1984

EQ 10GM BASE/VIAL N050579 004 May 23, 1984

CEFOPERAZONE SODIUM

INJECTABLE; INJECTION

CEFOBID

PFIZER EQ 1GM BASE/VIAL A063333 001 Mar 31, 1995

EQ 1GM BASE/VIAL N050551 001 Nov 18, 1982

EQ 2GM BASE/VIAL A063333 002 Mar 31, 1995

EQ 2GM BASE/VIAL N050551 002 Nov 18, 1982

EQ 10GM BASE/VIAL N050551 003 Mar 05, 1990

CEFOBID IN PLASTIC CONTAINER

PFIZER EQ 20MG BASE/ML N050613 002 Jul 31, 1987

EQ 40MG BASE/ML N050613 001 Jul 23, 1986

CEFORANIDE

INJECTABLE; INJECTION

PRECEF

APOTHECON 500MG/VIAL A062579 001 Nov 26, 1984

1GM/VIAL A062579 002 Nov 26, 1984

2GM/VIAL A062579 003 Nov 26, 1984

10GM/VIAL A062579 004 Nov 26, 1984

20GM/VIAL A062579 005 Nov 26, 1984

BRISTOL 500MG/VIAL N050554 001 May 24, 1984

1GM/VIAL N050554 002 May 24, 1984

2GM/VIAL N050554 003 May 24, 1984

10GM/VIAL N050554 004 May 24, 1984

20GM/VIAL N050554 005 May 24, 1984

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CEFOTAXIME SODIUM

INJECTABLE; INJECTION

CEFOTAXIME

FRESENIUS KABI USA	EQ 500MG BASE/VIAL	A064200 001	Mar 24, 2000
	EQ 1GM BASE/VIAL	A064200 002	Mar 24, 2000
	EQ 2GM BASE/VIAL	A064200 003	Mar 24, 2000
	EQ 10GM BASE/VIAL	A064201 001	Mar 24, 2000
	EQ 20GM BASE/VIAL	A064201 002	Mar 24, 2000
HIKMA	EQ 500MG BASE/VIAL	A065072 001	Nov 20, 2002
	EQ 1GM BASE/VIAL	A065072 002	Nov 20, 2002
	EQ 2GM BASE/VIAL	A065072 003	Nov 20, 2002
	EQ 10GM BASE/VIAL	A065071 001	Nov 20, 2002
WOCKHARDT	EQ 1GM BASE/VIAL	A065197 001	Aug 29, 2006
CEFOTAXIME AND DEXTROSE 2.4% IN PLASTIC CONTAINER			
B BRAUN	EQ 2GM BASE	N050792 001	Jul 29, 2004
CEFOTAXIME AND DEXTROSE 3.9% IN PLASTIC CONTAINER			
B BRAUN	EQ 1GM BASE	N050792 002	Jul 29, 2004
CEFOTAXIME SODIUM			
AUROBINDO PHARMA	EQ 500MG BASE/VIAL	A065517 001	Nov 06, 2009
	EQ 1GM BASE/VIAL	A065517 002	Nov 06, 2009
	EQ 2GM BASE/VIAL	A065517 003	Nov 06, 2009
AUROBINDO PHARMA LTD	EQ 10GM BASE/VIAL	A065516 001	Nov 06, 2009
CEPHAZONE PHARMA	EQ 10GM BASE/VIAL	A065348 001	Jan 25, 2010
HOSPIRA	EQ 1GM BASE/VIAL	A203132 001	Feb 19, 2016
	EQ 2GM BASE/VIAL	A203132 002	Feb 19, 2016
HOSPIRA INC	EQ 500MG BASE/VIAL	A065290 001	Aug 11, 2006
	EQ 1GM BASE/VIAL	A065290 002	Aug 11, 2006
	EQ 1GM BASE/VIAL	A065293 001	Aug 10, 2006
	EQ 2GM BASE/VIAL	A065290 003	Aug 11, 2006
	EQ 2GM BASE/VIAL	A065293 002	Aug 10, 2006
	EQ 10GM BASE/VIAL	A065292 001	Aug 10, 2006
LUPIN	EQ 500MG BASE/VIAL	A065124 001	Sep 24, 2003
	EQ 1GM BASE/VIAL	A065124 002	Sep 24, 2003
	EQ 2GM BASE/VIAL	A065124 003	Sep 24, 2003
WOCKHARDT	EQ 500MG BASE/VIAL	A065197 002	Jun 20, 2008
	EQ 2GM BASE/VIAL	A065197 003	Jun 20, 2008
CLAFORAN			
SANOFI AVENTIS US	EQ 1GM BASE/VIAL	A062659 001	Jan 13, 1987
	EQ 2GM BASE/VIAL	A062659 002	Jan 13, 1987
+ STERIMAX	EQ 500MG BASE/VIAL **	N050547 001	
+	EQ 1GM BASE/VIAL **	N050547 002	
+	EQ 2GM BASE/VIAL **	N050547 003	
+	EQ 10GM BASE/VIAL **	N050547 004	Dec 29, 1983
CLAFORAN IN DEXTROSE 5% IN PLASTIC CONTAINER			
STERIMAX	EQ 20MG BASE/ML	N050596 002	May 20, 1985
	EQ 40MG BASE/ML	N050596 004	May 20, 1985
CLAFORAN IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER			
STERIMAX	EQ 20MG BASE/ML	N050596 001	May 20, 1985
	EQ 40MG BASE/ML	N050596 003	May 20, 1985

CEFOTETAN DISODIUM

INJECTABLE; INJECTION

CEFOTAN

PAI HOLDINGS PHARM	EQ 10GM BASE/VIAL **	N050588 003	Apr 25, 1988
TELIGENT	EQ 1GM BASE/VIAL	A063293 001	Apr 29, 1993
	EQ 2GM BASE/VIAL	A063293 002	Apr 29, 1993
CEFOTAN IN PLASTIC CONTAINER			
PAI HOLDINGS PHARM	EQ 20MG BASE/ML	N050694 002	Jul 30, 1993
	EQ 40MG BASE/ML	N050694 001	Jul 30, 1993
CEFOTETAN			
FRESENIUS KABI USA	EQ 10GM BASE/VIAL	A065375 001	Aug 09, 2007
HIKMA	EQ 1GM BASE/VIAL	A091031 001	Oct 26, 2011
	EQ 2GM BASE/VIAL	A091031 002	Oct 26, 2011
WEST-WARD PHARM CORP	EQ 10GM BASE/VIAL	A091030 001	Oct 26, 2011
CEFOTETAN AND DEXTROSE IN DUPLEX CONTAINER			
+ B BRAUN	EQ 1GM BASE/VIAL	N065430 001	Aug 09, 2007
+	EQ 2GM BASE/VIAL	N065430 002	Aug 09, 2007

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CEFOTIAM HYDROCHLORIDE

INJECTABLE; INJECTION

CERADON

TAKEDA

EQ 1GM BASE/VIAL

N050601 001 Dec 30, 1988

CEFOXITIN SODIUM

INJECTABLE; INJECTION

CEFOXITIN

ACS DOBFAR SPA

EQ 1GM BASE/VIAL

A065467 001 Aug 31, 2011

EQ 2GM BASE/VIAL

A065467 002 Aug 31, 2011

EQ 10GM BASE/VIAL

A065464 001 Aug 31, 2011

FRESENIUS KABI USA

EQ 1GM BASE/VIAL **

A065012 001 Jul 03, 2000

EQ 2GM BASE/VIAL **

A065012 002 Jul 03, 2000

EQ 10GM BASE/VIAL

A065011 001 Jul 03, 2000

HOSPIRA INC

EQ 1GM BASE/VIAL

A065313 001 Jan 23, 2006

EQ 2GM BASE/VIAL

A065313 002 Jan 23, 2006

EQ 10GM BASE/VIAL

A065312 001 Feb 13, 2006

MEFOXIN

NORVIUM BIOSCIENCE

EQ 1GM BASE/VIAL

A062757 001 Jan 08, 1987

+

EQ 1GM BASE/VIAL **

N050517 001

EQ 2GM BASE/VIAL

A062757 002 Jan 08, 1987

+

EQ 2GM BASE/VIAL **

N050517 002

+

EQ 10GM BASE/VIAL **

N050517 003

MEFOXIN IN DEXTROSE 5% IN PLASTIC CONTAINER

+

MERCK

EQ 20MG BASE/ML **

N050581 003 Sep 20, 1984

+

EQ 40MG BASE/ML **

N050581 004 Sep 20, 1984

MEFOXIN IN PLASTIC CONTAINER

NORVIUM BIOSCIENCE

EQ 20MG BASE/ML

A063182 001 Jan 25, 1993

EQ 40MG BASE/ML

A063182 002 Jan 25, 1993

MEFOXIN IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

+

MERCK

EQ 20MG BASE/ML **

N050581 002 Sep 20, 1984

+

EQ 40MG BASE/ML **

N050581 001 Sep 20, 1984

CEFPYRAMIDE SODIUM

INJECTABLE; INJECTION

CEFPYRAMIDE SODIUM

WYETH AYERST

EQ 1GM BASE/VIAL

N050633 002 Jan 31, 1989

EQ 2GM BASE/VIAL

N050633 003 Jan 31, 1989

EQ 10GM BASE/VIAL

N050633 005 Jan 31, 1989

CEFPODOXIME PROXETIL

FOR SUSPENSION; ORAL

BANAN

SANKYO

EQ 50MG BASE/5ML

N050688 002 Aug 07, 1992

EQ 100MG BASE/5ML

N050688 001 Aug 07, 1992

CEFPODOXIME PROXETIL

CHARTWELL RX

EQ 50MG BASE/5ML

A090031 001 Jan 14, 2009

EQ 100MG BASE/5ML

A090031 002 Jan 14, 2009

SUN PHARM INDS LTD

EQ 50MG BASE/5ML

A065082 001 May 31, 2002

EQ 100MG BASE/5ML

A065082 002 May 31, 2002

VANTIN

+

PHARMACIA AND UPJOHN

EQ 50MG BASE/5ML **

N050675 001 Aug 07, 1992

+

EQ 100MG BASE/5ML **

N050675 002 Aug 07, 1992

TABLET; ORAL

BANAN

SANKYO

EQ 100MG BASE

N050687 001 Aug 07, 1992

EQ 200MG BASE

N050687 002 Aug 07, 1992

CEFPODOXIME PROXETIL

SUN PHARM INDS LTD

EQ 100MG BASE

A065083 001 Aug 20, 2003

EQ 200MG BASE

A065083 002 Aug 20, 2003

VANTIN

+

PHARMACIA AND UPJOHN

EQ 100MG BASE **

N050674 001 Aug 07, 1992

+

EQ 200MG BASE **

N050674 002 Aug 07, 1992

CEFPROZIL

FOR SUSPENSION; ORAL

CEFPROZIL

CHARTWELL RX

125MG/5ML

A065236 001 Dec 08, 2005

250MG/5ML

A065236 002 Dec 08, 2005

ORCHID HLTHCARE

125MG/5ML

A065284 002 Dec 30, 2005

250MG/5ML

A065284 001 Dec 30, 2005

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CEFPROZIL

FOR SUSPENSION; ORAL

CEFPROZIL

RANBAXY LABS LTD	125MG/5ML	A065202 001	Jun 30, 2006
	250MG/5ML	A065202 002	Jun 30, 2006
SANDOZ	125MG/5ML	A065257 001	Dec 08, 2005
	250MG/5ML	A065257 002	Dec 08, 2005

CEFZIL

+ CORDEN PHARMA	125MG/5ML **	N050665 001	Dec 23, 1991
+	250MG/5ML **	N050665 002	Dec 23, 1991

TABLET; ORAL

CEFPROZIL

ORCHID HLTHCARE	250MG	A065267 001	Dec 19, 2005
	500MG	A065267 002	Dec 19, 2005
RANBAXY LABS LTD	250MG	A065198 001	Dec 13, 2006
	500MG	A065198 002	Dec 13, 2006
WOCKHARDT	250MG	A065428 001	Jun 14, 2007
	500MG	A065428 002	Jun 14, 2007

CEFZIL

+ CORDEN PHARMA	250MG **	N050664 001	Dec 23, 1991
+	500MG **	N050664 002	Dec 23, 1991

CEFTAROLINE FOSAMIL

POWDER; INTRAVENOUS

CEFTAROLINE FOSAMIL

APOTEX	400MG/VIAL	A208075 001	Sep 21, 2021
	600MG/VIAL	A208075 002	Sep 21, 2021

CEFTAZIDIME

INJECTABLE; INJECTION

CEFTAZIDIME

ACS DOBFAR	500MG/VIAL	A062640 001	Nov 20, 1985
AUROBINDO PHARMA LTD	500MG/VIAL	A065481 001	May 28, 2010
	1GM/VIAL	A065481 002	May 28, 2010
	2GM/VIAL	A065481 003	May 28, 2010
	6GM/VIAL	A065482 001	May 28, 2010
WOCKHARDT	1GM/VIAL	A065196 001	Oct 15, 2008

CEFTAZIDIME IN DEXTROSE CONTAINER

+ B BRAUN	EQ 1GM BASE	N050823 001	Jun 13, 2011
+	EQ 2GM BASE	N050823 002	Jun 13, 2011

CEPTAZ

GLAXOSMITHKLINE	500MG/VIAL	N050646 001	Sep 27, 1990
	1GM/VIAL	N050646 002	Sep 27, 1990
	2GM/VIAL	N050646 003	Sep 27, 1990
	10GM/VIAL	N050646 004	Sep 27, 1990

FORTAZ

+ PAI HOLDINGS PHARM	500MG/VIAL **	N050578 001	Jul 19, 1985
+	1GM/VIAL **	N050578 002	Jul 19, 1985
+	2GM/VIAL **	N050578 003	Jul 19, 1985
+	6GM/VIAL **	N050578 004	Jul 19, 1985

PENTACEF

GLAXOSMITHKLINE	1GM/VIAL	A063322 001	Nov 07, 1995
	1GM/VIAL	A064006 001	Mar 31, 1992
	2GM/VIAL	A063322 002	Nov 07, 1995
	2GM/VIAL	A064006 002	Mar 31, 1992
	6GM/VIAL	A064008 001	Mar 31, 1992
	10GM/VIAL	A064008 002	Mar 31, 1992

TAZICEF

HOSPIRA	500MG/VIAL	A062662 001	Mar 06, 1986
---------	------------	-------------	--------------

TAZIDIME

LILLY	1GM/VIAL	A062655 001	Nov 20, 1985
	2GM/VIAL	A062655 002	Nov 20, 1985

TAZIDIME IN PLASTIC CONTAINER

LILLY	1GM/VIAL	A062739 001	Jul 10, 1986
	2GM/VIAL	A062739 002	Jul 10, 1986

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CEFTAZIDIME SODIUM

INJECTABLE; INJECTION

CEFTAZIDIME SODIUM IN PLASTIC CONTAINER

BAXTER HLTHCARE	EQ 10MG BASE/ML	A063221 001	Apr 29, 1993
	EQ 20MG BASE/ML	A063221 002	Apr 29, 1993
	EQ 40MG BASE/ML	A063221 003	Apr 29, 1993

FORTAZ IN PLASTIC CONTAINER

PAI HOLDINGS PHARM	EQ 10MG BASE/ML	N050634 001	Apr 28, 1989
+	EQ 20MG BASE/ML	N050634 002	Apr 28, 1989
+	EQ 40MG BASE/ML	N050634 003	Apr 28, 1989

CEFTIBUTEN DIHYDRATE

CAPSULE; ORAL

CEDAX

SI PHARMS	EQ 400MG BASE	N050685 002	Dec 20, 1995
-----------	---------------	-------------	--------------

FOR SUSPENSION; ORAL

CEDAX

+	SI PHARMS	EQ 90MG BASE/5ML **	N050686 001	Dec 20, 1995
+		EQ 180MG BASE/5ML **	N050686 002	Dec 20, 1995

CEFTIZOXIME SODIUM

INJECTABLE; INJECTION

CEFIZOX

ASTELLAS	EQ 500MG BASE/VIAL	N050560 001	Sep 15, 1983
	EQ 1GM BASE/VIAL	A063294 002	Mar 31, 1994
	EQ 1GM BASE/VIAL	N050560 002	Sep 15, 1983
	EQ 2GM BASE/VIAL	A063294 003	Mar 31, 1994
	EQ 2GM BASE/VIAL	N050560 003	Sep 15, 1983
	EQ 10GM BASE/VIAL	N050560 005	Mar 19, 1993

CEFIZOX IN DEXTROSE 5% IN PLASTIC CONTAINER

ASTELLAS	EQ 20MG BASE/ML	N050589 001	Oct 03, 1984
	EQ 40MG BASE/ML	N050589 002	Oct 03, 1984

CEFIZOX IN PLASTIC CONTAINER

ASTELLAS	EQ 20MG BASE/ML	N050589 003	Apr 13, 1995
	EQ 40MG BASE/ML	N050589 004	Apr 13, 1995

CEFTOBIPOLE MEDOCARIL SODIUM

POWDER; INTRAVENOUS

ZEVTERA

+	BASILEA PHARM ALLSCH	667MG/VIAL	N218275 001	Apr 03, 2024
---	----------------------	------------	-------------	--------------

CEFTRIAZONE SODIUM

INJECTABLE; INJECTION

CEFTRIAZONE

AGILA SPECLTS	EQ 10GM BASE/VIAL	A091068 001	Jan 07, 2013
AUROBINDO PHARMA LTD	EQ 10GM BASE/VIAL	A065504 001	Jul 31, 2008
BEDFORD	EQ 10GM BASE/VIAL	A065475 001	Aug 18, 2008
FACTA FARMA	EQ 10GM BASE/VIAL	A065269 001	Feb 28, 2007
FRESENIUS KABI USA	EQ 10GM BASE/VIAL	A065252 001	Feb 15, 2006
HOSPIRA INC	EQ 1GM BASE/VIAL	A065231 001	Aug 02, 2005
	EQ 1GM BASE/VIAL	A202563 001	Aug 20, 2012
	EQ 2GM BASE/VIAL	A065231 002	Aug 02, 2005
	EQ 2GM BASE/VIAL	A202563 002	Aug 20, 2012
	EQ 10GM BASE/VIAL	A065232 001	Aug 02, 2005
LUPIN	EQ 10GM BASE/VIAL	A065263 001	Sep 12, 2006
TEVA	EQ 10GM BASE/VIAL	A065274 001	May 01, 2006

ROCEPHIN

HOFFMANN LA ROCHE	EQ 250MG BASE/VIAL	A063239 001	Aug 13, 1993
	EQ 500MG BASE/VIAL	A062654 001	Apr 30, 1987
	EQ 500MG BASE/VIAL	A063239 002	Aug 13, 1993
	EQ 1GM BASE/VIAL	A062654 002	Apr 30, 1987
	EQ 1GM BASE/VIAL	A063239 003	Aug 13, 1993
	EQ 2GM BASE/VIAL	A062654 003	Apr 30, 1987
+	EQ 10GM BASE/VIAL **	N050585 005	Dec 21, 1984
ROCHE	EQ 250MG BASE/VIAL	A062510 001	Mar 12, 1985
	EQ 500MG BASE/VIAL	A062510 002	Mar 12, 1985
	EQ 1GM BASE/VIAL	A062510 003	Mar 12, 1985

ROCEPHIN W/ DEXTROSE IN PLASTIC CONTAINER

+	HOFFMANN LA ROCHE	EQ 10MG BASE/ML **	N050624 001	Feb 11, 1987
+		EQ 20MG BASE/ML **	N050624 002	Feb 11, 1987
+		EQ 40MG BASE/ML **	N050624 003	Feb 11, 1987

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CEFTRIAZONE SODIUM

INJECTABLE; INTRAMUSCULAR, INTRAVENOUS

CEFTRIAZONE

ASTRAL	EQ 250MG BASE/VIAL	A091049 001	Jun 11, 2018
	EQ 500MG BASE/VIAL	A091049 002	Jun 11, 2018
	EQ 1GM BASE/VIAL	A091049 003	Jun 11, 2018
	EQ 2GM BASE/VIAL	A091049 004	Jun 11, 2018
AUROBINDO PHARMA LTD	EQ 250MG BASE/VIAL	A065505 001	Jul 31, 2008
	EQ 500MG BASE/VIAL	A065505 002	Jul 31, 2008
	EQ 1GM BASE/VIAL	A065505 003	Jul 31, 2008
	EQ 2GM BASE/VIAL	A065505 004	Jul 31, 2008
BEDFORD	EQ 250MG BASE/VIAL	A065465 001	Aug 18, 2008
	EQ 500MG BASE/VIAL	A065465 002	Aug 18, 2008
	EQ 1GM BASE/VIAL	A065465 003	Aug 18, 2008
	EQ 2GM BASE/VIAL	A065465 004	Aug 18, 2008
CEPHAZONE PHARMA	EQ 250MG BASE/VIAL	A065294 001	Mar 26, 2007
	EQ 500MG BASE/VIAL	A065294 002	Mar 26, 2007
	EQ 1GM BASE/VIAL	A065294 003	Mar 26, 2007
	EQ 2GM BASE/VIAL	A065294 004	Mar 26, 2007
DEVA HOLDING AS	EQ 250MG BASE/VIAL	A210197 001	Jan 12, 2024
	EQ 500MG BASE/VIAL	A210197 002	Jan 12, 2024
	EQ 1GM BASE/VIAL	A210197 003	Jan 12, 2024
	EQ 2GM BASE/VIAL	A210197 004	Jan 12, 2024
FACTA FARMA	EQ 1GM BASE/VIAL	A065268 001	Feb 28, 2007
	EQ 2GM BASE/VIAL	A065268 002	Feb 28, 2007
FRESENIUS KABI USA	EQ 250MG BASE/VIAL	A065245 001	Feb 15, 2006
	EQ 500MG BASE/VIAL	A065245 002	Feb 15, 2006
	EQ 1GM BASE/VIAL	A065245 003	Feb 15, 2006
	EQ 2GM BASE/VIAL	A065245 004	Feb 15, 2006
HOSPIRA INC	EQ 250MG BASE/VIAL	A065230 001	Aug 02, 2005
	EQ 500MG BASE/VIAL	A065230 002	Aug 02, 2005
	EQ 1GM BASE/VIAL	A065230 003	Aug 02, 2005
	EQ 2GM BASE/VIAL	A065230 004	Aug 02, 2005
LUPIN	EQ 250MG BASE/VIAL	A065125 001	Sep 30, 2003
	EQ 500MG BASE/VIAL	A065125 002	Sep 30, 2003
	EQ 1GM BASE/VIAL	A065125 003	Sep 30, 2003
	EQ 2GM BASE/VIAL	A065125 004	Sep 30, 2003
TEVA	EQ 1GM BASE/VIAL	A065262 001	Jun 29, 2006
	EQ 2GM BASE/VIAL	A065262 002	Jun 29, 2006
TEVA PHARMS USA	EQ 250MG BASE/VIAL	A065227 001	Mar 15, 2007
	EQ 500MG BASE/VIAL	A065227 002	Mar 15, 2007
	EQ 1GM BASE/VIAL	A065227 003	Mar 15, 2007
	EQ 2GM BASE/VIAL	A065227 004	Mar 15, 2007
ROCEPHIN			
+ HOFFMANN LA ROCHE	EQ 250MG BASE/VIAL **	N050585 001	Dec 21, 1984
+ HOFFMANN LA ROCHE	EQ 500MG BASE/VIAL **	N050585 002	Dec 21, 1984
+ HOFFMANN LA ROCHE	EQ 1GM BASE/VIAL **	N050585 003	Dec 21, 1984
+ HOFFMANN LA ROCHE	EQ 2GM BASE/VIAL **	N050585 004	Dec 21, 1984

CEFTRIAZONE SODIUM; LIDOCAINE

INJECTABLE; INJECTION

ROCEPHIN KIT

HOFFMANN LA ROCHE	EQ 500MG BASE/VIAL, N/A; N/A, 1%	N050585 007	May 08, 1996
HOFFMANN LA ROCHE	EQ 1GM BASE/VIAL, N/A; N/A, 1%	N050585 006	May 08, 1996

CEFUROXIME AXETIL

FOR SUSPENSION; ORAL

CEFTIN

+ GLAXOSMITHKLINE	EQ 125MG BASE/5ML	N050672 001	Jun 30, 1994
+ GLAXOSMITHKLINE	EQ 250MG BASE/5ML	N050672 002	Apr 29, 1997

CEFUROXIME AXETIL

SUN PHARM INDS LTD	EQ 125MG BASE/5ML	A065323 001	Feb 05, 2008
SUN PHARM INDS LTD	EQ 250MG BASE/5ML	A065323 002	Feb 05, 2008

TABLET; ORAL

CEFTIN

+ GLAXOSMITHKLINE	EQ 125MG BASE **	N050605 001	Dec 28, 1987
+ GLAXOSMITHKLINE	EQ 250MG BASE **	N050605 002	Dec 28, 1987
+ GLAXOSMITHKLINE	EQ 500MG BASE **	N050605 003	Dec 28, 1987

CEFUROXIME AXETIL

ANI PHARMS	EQ 250MG BASE	A065190 001	Oct 18, 2004
------------	---------------	-------------	--------------

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CEFUROXIME AXETIL

TABLET; ORAL

CEFUROXIME AXETIL

	EQ 500MG BASE	A065190 002	Oct 18, 2004
FOSUN PHARMA	EQ 250MG BASE	A065126 001	Oct 28, 2003
	EQ 500MG BASE	A065126 002	Oct 28, 2003
RANBAXY LABS LTD	EQ 125MG BASE	A065043 003	Feb 15, 2002
	EQ 250MG BASE	A065043 002	Feb 15, 2002
	EQ 500MG BASE	A065043 001	Feb 15, 2002
SUN PHARM INDS LTD	EQ 125MG BASE	A065118 001	Apr 25, 2003
	EQ 250MG BASE	A065118 002	Apr 25, 2003
	EQ 500MG BASE	A065118 003	Apr 25, 2003

CEFUROXIME SODIUM

INJECTABLE; INJECTION

CEFUROXIME AND DEXTROSE IN DUPLEX CONTAINER

+ B BRAUN	EQ 750MG BASE/VIAL	N050780 001	Feb 21, 2001
-----------	--------------------	-------------	--------------

+	EQ 1.5GM BASE/VIAL	N050780 002	Feb 21, 2001
---	--------------------	-------------	--------------

CEFUROXIME SODIUM

ACS DOBFAR SPA	EQ 7.5GM BASE/VIAL	A064124 001	May 30, 1997
----------------	--------------------	-------------	--------------

FRESENIUS KABI USA	EQ 1.5GM BASE/VIAL	A065001 002	May 30, 2001
--------------------	--------------------	-------------	--------------

	EQ 7.5GM BASE/VIAL	A065002 001	Sep 28, 1998
--	--------------------	-------------	--------------

HIKMA	EQ 7.5GM BASE/VIAL	A065046 001	Jan 09, 2004
-------	--------------------	-------------	--------------

HOSPIRA INC	EQ 1.5GM BASE/VIAL	A065483 002	Oct 15, 2008
-------------	--------------------	-------------	--------------

	EQ 1.5GM BASE/VIAL	A065503 001	Oct 15, 2008
--	--------------------	-------------	--------------

	EQ 7.5GM BASE/VIAL	A065484 001	Oct 15, 2008
--	--------------------	-------------	--------------

TEVA PHARMS	EQ 7.5GM BASE/VIAL	A064191 001	Apr 16, 1998
-------------	--------------------	-------------	--------------

WATSON LABS INC	EQ 1.5GM BASE/VIAL	A064035 002	Feb 26, 1993
-----------------	--------------------	-------------	--------------

	EQ 7.5GM BASE/VIAL	A064036 001	Feb 26, 1993
--	--------------------	-------------	--------------

CEFUROXIME SODIUM IN PLASTIC CONTAINER

SAMSON MEDCL	EQ 75GM BASE/VIAL	A065251 001	Dec 30, 2009
--------------	-------------------	-------------	--------------

	EQ 225GM BASE/VIAL	A065251 002	Dec 30, 2009
--	--------------------	-------------	--------------

KEFUROX

ACS DOBFAR	EQ 1.5GM BASE/VIAL	A062591 002	Jan 10, 1986
------------	--------------------	-------------	--------------

	EQ 7.5GM BASE/VIAL	A062591 003	Dec 17, 1987
--	--------------------	-------------	--------------

LILLY	EQ 1.5GM BASE/VIAL	A062592 002	Jan 10, 1986
-------	--------------------	-------------	--------------

KEFUROX IN PLASTIC CONTAINER

LILLY	EQ 1.5GM BASE/VIAL	A062590 002	Jan 10, 1986
-------	--------------------	-------------	--------------

ZINACEF

+ PAI HOLDINGS PHARM	EQ 1.5GM BASE/VIAL **	N050558 003	Oct 19, 1983
----------------------	-----------------------	-------------	--------------

+	EQ 7.5GM BASE/VIAL **	N050558 004	Oct 23, 1986
---	-----------------------	-------------	--------------

ZINACEF IN PLASTIC CONTAINER

PAI HOLDINGS PHARM	EQ 15MG BASE/ML	N050643 001	Apr 28, 1989
--------------------	-----------------	-------------	--------------

+	EQ 30MG BASE/ML	N050643 002	Apr 28, 1989
---	-----------------	-------------	--------------

INJECTABLE; INTRAMUSCULAR, INTRAVENOUS

CEFUROXIME SODIUM

FRESENIUS KABI USA	EQ 750MG BASE/VIAL	A065001 001	May 30, 2001
--------------------	--------------------	-------------	--------------

HOSPIRA INC	EQ 750MG BASE/VIAL	A065483 001	Oct 15, 2008
-------------	--------------------	-------------	--------------

TEVA PHARMS	EQ 750MG BASE/VIAL	A064192 002	Apr 16, 1998
-------------	--------------------	-------------	--------------

	EQ 1.5GM BASE/VIAL	A064192 001	Apr 16, 1998
--	--------------------	-------------	--------------

WATSON LABS INC	EQ 750MG BASE/VIAL	A064035 001	Feb 26, 1993
-----------------	--------------------	-------------	--------------

KEFUROX

ACS DOBFAR	EQ 750MG BASE/VIAL	A062591 001	Jan 10, 1986
------------	--------------------	-------------	--------------

ZINACEF

+ PAI HOLDINGS PHARM	EQ 750MG BASE/VIAL **	N050558 002	Oct 19, 1983
----------------------	-----------------------	-------------	--------------

INJECTABLE; INTRAVENOUS

KEFUROX

LILLY	EQ 750MG BASE/VIAL	A062592 001	Jan 10, 1986
-------	--------------------	-------------	--------------

KEFUROX IN PLASTIC CONTAINER

LILLY	EQ 750MG BASE/VIAL	A062590 001	Jan 10, 1986
-------	--------------------	-------------	--------------

CELECOXIB

CAPSULE; ORAL

CELECOXIB

ACIC PHARMS	200MG	A212925 001	Dec 09, 2020
-------------	-------	-------------	--------------

	400MG	A212925 002	Dec 09, 2020
--	-------	-------------	--------------

AMNEAL PHARMS	50MG	A208833 001	May 31, 2018
---------------	------	-------------	--------------

	100MG	A208833 002	May 31, 2018
--	-------	-------------	--------------

	200MG	A208833 003	May 31, 2018
--	-------	-------------	--------------

	400MG	A208833 004	May 31, 2018
--	-------	-------------	--------------

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CELECOXIB

CAPSULE; ORAL

CELECOXIB

JUBILANT GENERICS	50MG	A207061 001	Apr 04, 2017
	100MG	A207061 002	Apr 04, 2017
	200MG	A207061 003	Apr 04, 2017
	400MG	A207061 004	Apr 04, 2017
NORVIUM BIOSCIENCE	50MG	A078857 001	May 30, 2014
	100MG	A078857 002	Feb 11, 2015
	200MG	A078857 003	Feb 11, 2015
	400MG	A078857 004	Feb 11, 2015
UNICHEM	50MG	A213301 001	Jan 12, 2021
	100MG	A213301 002	Jan 12, 2021
	200MG	A213301 003	Jan 12, 2021
	400MG	A213301 004	Jan 12, 2021
YABAO PHARM	100MG	A212564 001	Apr 10, 2023
	200MG	A212564 002	Apr 10, 2023
	400MG	A212564 003	Apr 10, 2023

CELECOXIB; TRAMADOL HYDROCHLORIDE

TABLET; ORAL

SEGLENTIS

+ KOWA PHARMS	56MG; 44MG	N213426 001	Oct 15, 2021
---------------	------------	-------------	--------------

CELLULOSE SODIUM PHOSPHATE

POWDER; ORAL

CALCIBIND

MISSION PHARMA	2.5GM/PACKET	N018757 002	Dec 28, 1982
	300GM/BOT	N018757 003	Oct 16, 1984

CEPHALEXIN

CAPSULE; ORAL

CEPHALEXIN

APOTHECON	EQ 250MG BASE	A062973 001	Nov 08, 1988
	EQ 250MG BASE	A063063 001	Sep 29, 1989
	EQ 250MG BASE	A063186 001	Dec 30, 1994
	EQ 500MG BASE	A062974 001	Nov 23, 1988
	EQ 500MG BASE	A063063 002	Sep 29, 1989
	EQ 500MG BASE	A063186 002	Dec 30, 1994
BARR	EQ 250MG BASE	A062773 001	Jun 26, 1987
	EQ 500MG BASE	A062775 001	Apr 22, 1987
FACTA FARMA	EQ 250MG BASE	A062118 001	
	EQ 500MG BASE	A062118 002	
HIKMA	EQ 250MG BASE	A065215 001	Jan 24, 2006
	EQ 500MG BASE	A065215 002	Jan 24, 2006
IVAX SUB TEVA PHARMS	EQ 250MG BASE	A061969 001	
	EQ 500MG BASE	A061969 002	
PUREPAC PHARM	EQ 250MG BASE	A062809 001	Apr 22, 1987
	EQ 500MG BASE	A062809 002	Apr 22, 1987
STEVENS J	EQ 250MG BASE	A062870 001	Mar 17, 1988
	EQ 500MG BASE	A062869 001	Mar 17, 1988
SUN PHARM INDS (IN)	EQ 250MG BASE	A062791 001	Jun 11, 1987
	EQ 500MG BASE	A062791 002	Jun 11, 1987
SUN PHARM INDS LTD	EQ 250MG BASE	A065007 001	Sep 16, 1999
	EQ 500MG BASE	A065007 002	Sep 16, 1999
TEVA	EQ 250MG BASE	A062760 001	Apr 24, 1987
	EQ 250MG BASE	A062821 001	Feb 05, 1988
	EQ 500MG BASE	A062761 001	Apr 24, 1987
	EQ 500MG BASE	A062823 001	Feb 05, 1988
YOSHITOMI	EQ 250MG BASE	A062872 001	Jun 20, 1988
	EQ 500MG BASE	A062871 001	Jul 05, 1988
KEFLEX			
+ PRAGMA	EQ 250MG BASE **	N050405 002	
+	EQ 333MG BASE **	N050405 004	May 12, 2006
+	EQ 500MG BASE **	N050405 003	
+	EQ 750MG BASE **	N050405 005	May 12, 2006

FOR SUSPENSION; ORAL

CEPHALEXIN

APOTHECON	EQ 125MG BASE/5ML	A062986 001	Apr 18, 1991
	EQ 250MG BASE/5ML	A062987 001	Jul 25, 1989
BARR	EQ 125MG BASE/5ML	A062778 001	Aug 06, 1987

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CEPHALEXINFOR SUSPENSION;ORAL
CEPHALEXIN

	EQ 250MG BASE/5ML	A062777 001	Aug 06, 1987
CHARTWELL RX	EQ 125MG BASE/5ML	A065326 001	Jul 10, 2006
	EQ 250MG BASE/5ML	A065326 002	Jul 10, 2006
FACTA FARMA	EQ 100MG BASE/ML **	A062117 001	
	EQ 125MG BASE/5ML **	A062117 002	
	EQ 250MG BASE/5ML **	A062117 003	
HIKMA PHARMS	EQ 125MG BASE/5ML	A065444 001	Aug 28, 2009
	EQ 250MG BASE/5ML	A065444 002	Aug 28, 2009
SUN PHARM INDS LTD	EQ 125MG BASE/5ML	A065081 001	Jul 27, 2001
	EQ 250MG BASE/5ML	A065081 002	Jul 27, 2001
TEVA	EQ 125MG BASE/5ML	A062767 001	Jun 16, 1987
	EQ 125MG BASE/5ML	A062873 001	May 23, 1988
	EQ 250MG BASE/5ML	A062768 001	Jun 16, 1987
	EQ 250MG BASE/5ML	A062867 001	Apr 15, 1988
VITARINE	EQ 125MG BASE/5ML	A062779 001	Dec 22, 1987
	EQ 250MG BASE/5ML	A062781 001	Dec 22, 1987
KEFLEX			
+ PRAGMA	EQ 100MG BASE/ML **	N050406 003	
+	EQ 125MG BASE/5ML **	N050406 001	
+	EQ 250MG BASE/5ML **	N050406 002	

TABLET;ORAL

CEPHALEXIN

BARR	EQ 250MG BASE	A062826 001	Aug 17, 1987
	EQ 500MG BASE	A062827 001	Aug 17, 1987
VITARINE	EQ 250MG BASE	A062863 001	Aug 11, 1988
	EQ 500MG BASE	A062863 002	Aug 11, 1988
	EQ 1GM BASE	A062863 003	Aug 11, 1988
KEFLET			
LILLY	EQ 250MG BASE	A062745 001	Dec 01, 1986
+	EQ 250MG BASE **	N050440 003	Feb 26, 1987
	EQ 500MG BASE	A062745 002	Dec 01, 1986
+	EQ 500MG BASE **	N050440 001	
	EQ 1GM BASE	N050440 002	
TABLET, FOR SUSPENSION;ORAL			
PANIXINE DISPERDOSE			
RANBAXY LABS LTD	EQ 125MG BASE	A065100 002	Sep 11, 2003
	EQ 250MG BASE	A065100 001	Sep 11, 2003

CEPHALEXIN HYDROCHLORIDE

TABLET;ORAL

KEFTAB

LILLY	EQ 250MG BASE	N050614 001	Oct 29, 1987
	EQ 333MG BASE	N050614 003	May 16, 1988
	EQ 500MG BASE	N050614 002	Oct 29, 1987

CEPHALOGLYCIN

CAPSULE;ORAL

KAFOCIN

LILLY	250MG	N050219 001	
-------	-------	-------------	--

CEPHALOTHIN SODIUM

INJECTABLE;INJECTION

CEPHALOTHIN

INTL MEDICATION	EQ 500MG BASE/VIAL	A062426 001	May 03, 1985
	EQ 1GM BASE/VIAL	A062426 002	May 03, 1985
	EQ 2GM BASE/VIAL	A062426 003	May 03, 1985
	EQ 4GM BASE/VIAL	A062426 004	May 03, 1985
CEPHALOTHIN SODIUM			
ABBOTT	EQ 1GM BASE/VIAL	A062547 001	Sep 11, 1985
	EQ 1GM BASE/VIAL	A062548 001	Sep 11, 1985
	EQ 2GM BASE/VIAL	A062547 002	Sep 11, 1985
	EQ 2GM BASE/VIAL	A062548 002	Sep 11, 1985
ABRAXIS PHARM	EQ 1GM BASE/VIAL	A062666 002	Jun 10, 1987
	EQ 2GM BASE/VIAL	A062666 001	Jun 10, 1987
BRISTOL	EQ 1GM BASE/VIAL	A062464 001	May 07, 1984
	EQ 2GM BASE/VIAL	A062464 002	May 07, 1984
	EQ 4GM BASE/VIAL	A062464 003	May 07, 1984

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CEPHALOTHIN SODIUM

INJECTABLE; INJECTION

CEPHALOTHIN SODIUM W/ DEXTROSE IN PLASTIC CONTAINER

BAXTER HLTHCARE	EQ 20MG BASE/ML	A062422 003	Jan 31, 1984
	EQ 20MG BASE/ML	A062422 005	Jul 16, 1991
	EQ 20MG BASE/ML	A062730 001	Mar 05, 1987
	EQ 40MG BASE/ML	A062422 004	Jan 31, 1984
	EQ 40MG BASE/ML	A062422 006	Jul 16, 1991
	EQ 40MG BASE/ML	A062730 002	Mar 05, 1987

CEPHALOTHIN SODIUM W/ SODIUM CHLORIDE IN PLASTIC CONTAINER

BAXTER HLTHCARE	EQ 20MG BASE/ML	A062422 001	Jan 31, 1984
	EQ 40MG BASE/ML	A062422 002	Jan 31, 1984

KEFLIN

LILLY	EQ 1GM BASE/VIAL	N050482 001	
	EQ 2GM BASE/VIAL	N050482 002	
	EQ 4GM BASE/VIAL	N050482 003	
	EQ 20GM BASE/VIAL	N050482 007	

KEFLIN IN PLASTIC CONTAINER

LILLY	EQ 1GM BASE/VIAL	A062549 001	Sep 10, 1985
	EQ 2GM BASE/VIAL	A062549 002	Sep 10, 1985

SEFFIN

GLAXOSMITHKLINE	EQ 1GM BASE/VIAL	A062435 001	Nov 15, 1983
	EQ 2GM BASE/VIAL	A062435 002	Nov 15, 1983
	EQ 10GM BASE/VIAL	A062435 003	Nov 15, 1983

CEPHAPIRIN SODIUM

INJECTABLE; INJECTION

CEFADYL

APOTHECON	EQ 500MG BASE/VIAL	A062961 001	Sep 20, 1988
	EQ 500MG BASE/VIAL	N050446 005	
	EQ 1GM BASE/VIAL	A061769 001	
	EQ 1GM BASE/VIAL	A062724 001	Dec 23, 1986
	EQ 1GM BASE/VIAL	A062961 002	Sep 20, 1988
	EQ 1GM BASE/VIAL	N050446 001	
	EQ 2GM BASE/VIAL	A061769 002	
	EQ 2GM BASE/VIAL	A062724 002	Dec 23, 1986
	EQ 2GM BASE/VIAL	A062961 003	Sep 20, 1988
	EQ 2GM BASE/VIAL	N050446 002	
	EQ 4GM BASE/VIAL	A061769 003	
	EQ 4GM BASE/VIAL	A062961 004	Sep 20, 1988
	EQ 4GM BASE/VIAL	N050446 003	
	EQ 20GM BASE/VIAL	N050446 004	

CEPHAPIRIN SODIUM

ABRAXIS PHARM	EQ 500MG BASE/VIAL	A062723 001	Nov 17, 1986
	EQ 1GM BASE/VIAL	A062723 002	Nov 17, 1986
	EQ 2GM BASE/VIAL	A062723 003	Nov 17, 1986
	EQ 4GM BASE/VIAL	A062723 004	Nov 17, 1986
	EQ 20GM BASE/VIAL	A062723 005	Nov 17, 1986
HIKMA	EQ 500MG BASE/VIAL	A062720 001	Jul 02, 1987
	EQ 1GM BASE/VIAL	A062720 002	Jul 02, 1987
	EQ 2GM BASE/VIAL	A062720 003	Jul 02, 1987
	EQ 20GM BASE/VIAL	A062720 004	Jul 02, 1987

CEPHRADINE

CAPSULE; ORAL

ANSPOR

GLAXOSMITHKLINE	250MG	A061859 001	
	500MG	A061859 002	

CEPHRADINE

BARR	250MG	A062850 001	Apr 22, 1988
	500MG	A062851 001	Apr 22, 1988
IVAX SUB TEVA PHARMS	250MG	A062762 001	Mar 06, 1987
	500MG	A062762 002	Mar 06, 1987
TEVA	250MG	A062683 001	Jan 09, 1987
	500MG	A062683 002	Jan 09, 1987
VITARINE	250MG	A062813 001	Feb 25, 1988
	500MG	A062813 002	Feb 25, 1988

VELOSEF

APOTHECON	250MG	A061764 001	
	500MG	A061764 002	

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CEPHRADINE

CAPSULE; ORAL

VELOSEF '250'

ERSANA

250MG

N050548 001

VELOSEF '500'

ERSANA

500MG

N050548 002

FOR SUSPENSION; ORAL

ANSPOR

GLAXOSMITHKLINE

125MG/5ML

A061866 001

250MG/5ML

A061866 002

CEPHRADINE

BARR

125MG/5ML

A062858 001 May 19, 1988

250MG/5ML

A062859 001 May 19, 1988

TEVA

125MG/5ML

A062693 001 Jan 09, 1987

250MG/5ML

A062693 002 Jan 09, 1987

VELOSEF '125'

APOTHECON

125MG/5ML

A061763 001

VELOSEF '250'

APOTHECON

250MG/5ML

A061763 002

INJECTABLE; INJECTION

VELOSEF

APOTHECON

250MG/VIAL

A061976 001

500MG/VIAL

A061976 002

1GM/VIAL

A061976 004

2GM/VIAL

A061976 003

4GM/VIAL

A061976 005

TABLET; ORAL

VELOSEF

BRISTOL MYERS SQUIBB 1GM

N050530 001

CERITINIB

CAPSULE; ORAL

ZYKADIA

+ NOVARTIS

150MG

N205755 001 Apr 29, 2014

CERIVASTATIN SODIUM

TABLET; ORAL

BAYCOL

BAYER PHARMS

0.05MG

N020740 001 Jun 26, 1997

0.1MG

N020740 002 Jun 26, 1997

0.2MG

N020740 003 Jun 26, 1997

0.3MG

N020740 004 Jun 26, 1997

0.4MG

N020740 005 May 24, 1999

0.8MG

N020740 006 Jul 24, 2000

CERULETIDE DIETHYLAMINE

INJECTABLE; INJECTION

TYMTRAN

PHARMACIA AND UPJOHN 0.02MG/ML

N018296 001

CETIRIZINE HYDROCHLORIDE

CAPSULE; ORAL

CETIRIZINE HYDROCHLORIDE ALLERGY

APOTEX

10MG

A207235 001 Aug 12, 2016

SOLUTION; ORAL

CETIRIZINE HYDROCHLORIDE

ACTAVIS MID ATLANTIC 5MG/5ML

A078617 001 Feb 02, 2010

AUROBINDO PHARMA LTD 5MG/5ML

A090751 001 Dec 16, 2009

BIONPHARMA 5MG/5ML

A078488 001 Oct 06, 2008

LANNETT CO INC 5MG/5ML

A078496 001 Sep 25, 2009

RANBAXY LABS LTD 5MG/5ML

A077472 001 Jun 18, 2008

TEVA PHARMS 5MG/5ML

A077279 001 May 27, 2008

WOCKHARDT 5MG/5ML

A078757 001 Aug 28, 2009

CHILDREN'S CETIRIZINE HYDROCHLORIDE ALLERGY

CHARTWELL RX 5MG/5ML

A090378 002 May 09, 2008

CYPRESS PHARM 5MG/5ML

A090300 001 Oct 10, 2008

PHARM ASSOC 5MG/5ML

A090188 002 Apr 22, 2008

RANBAXY LABS LTD 5MG/5ML

A090183 002 Apr 24, 2008

TRIS PHARMA INC 5MG/5ML

A090572 001 Nov 16, 2012

CHILDREN'S CETIRIZINE HYDROCHLORIDE HIVES RELIEF

CHARTWELL RX 5MG/5ML

A090378 001 May 09, 2008

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CETIRIZINE HYDROCHLORIDE

SOLUTION;ORAL

CHILDREN'S CETIRIZINE HYDROCHLORIDE HIVES RELIEF

CYPRESS PHARM	5MG/5ML	A090300	002	Oct 10, 2008
PHARM ASSOC	5MG/5ML	A090188	001	Apr 22, 2008
RANBAXY LABS LTD	5MG/5ML	A090183	001	Apr 24, 2008
TRIS PHARMA INC	5MG/5ML	A090572	002	Nov 16, 2012

ZYRTEC

J AND J CONSUMER INC	5MG/5ML **	N020346	001	Sep 27, 1996
----------------------	------------	---------	-----	--------------

TABLET;ORAL

CETIRIZINE HYDROCHLORIDE ALLERGY

CIPLA LTD	5MG	A077318	001	Jul 25, 2013
	10MG	A077318	002	Jul 25, 2013
HERITAGE PHARMA	5MG	A078615	003	Dec 28, 2007
	10MG	A078615	004	Dec 28, 2007
IPCA LABS LTD	5MG	A202277	002	Mar 11, 2014
	10MG	A202277	004	Mar 11, 2014
SUN PHARM INDS INC	5MG	A077499	001	Dec 27, 2007
	10MG	A077499	002	Dec 27, 2007
TORRENT PHARMS LLC	5MG	A079191	001	Apr 15, 2010
	10MG	A079191	004	Apr 15, 2010
UNICHEM	5MG	A078680	003	Jun 26, 2009
	10MG	A078680	004	Jun 26, 2009

CETIRIZINE HYDROCHLORIDE HIVES

IPCA LABS LTD	5MG	A202277	001	Mar 11, 2014
	10MG	A202277	003	Mar 11, 2014
SUN PHARM INDS INC	5MG	A077499	003	Dec 27, 2007
	10MG	A077499	004	Dec 27, 2007
UNICHEM	5MG	A078680	001	Jun 26, 2009
	10MG	A078680	002	Jun 26, 2009

CETIRIZINE HYDROCHLORIDE HIVES RELIEF

TORRENT PHARMS LLC	5MG	A079191	003	Apr 15, 2010
	10MG	A079191	002	Apr 15, 2010

ZYRTEC ALLERGY

+ KENVUE BRANDS	5MG	N019835	003	Nov 16, 2007
-----------------	-----	---------	-----	--------------

ZYRTEC HIVES

+ KENVUE BRANDS	5MG	N019835	005	Nov 16, 2007
+ KENVUE BRANDS	10MG	N019835	006	Nov 16, 2007

TABLET, CHEWABLE;ORAL

CETIRIZINE HYDROCHLORIDE ALLERGY

SUN PHARM INDS INC	5MG	A077631	004	Jan 11, 2008
	10MG	A077631	003	Jan 11, 2008

CETIRIZINE HYDROCHLORIDE HIVES RELIEF

SUN PHARM INDS INC	5MG	A077631	001	Jan 11, 2008
	10MG	A077631	002	Jan 11, 2008

CHILDREN'S CETIRIZINE HYDROCHLORIDE ALLERGY

SUN PHARM	5MG	A090142	001	Aug 30, 2011
	10MG	A090142	002	Aug 30, 2011

CHILDREN'S CETIRIZINE HYDROCHLORIDE HIVES RELIEF

SUN PHARM	5MG	A090142	003	Aug 30, 2011
	10MG	A090142	004	Aug 30, 2011

CHILDREN'S ZYRTEC ALLERGY

+ KENVUE BRANDS	5MG **	N021621	003	Nov 16, 2007
-----------------	--------	---------	-----	--------------

CHILDREN'S ZYRTEC HIVES RELIEF

+ KENVUE BRANDS	5MG **	N021621	005	Nov 16, 2007
+ KENVUE BRANDS	10MG **	N021621	006	Nov 16, 2007

CETIRIZINE HYDROCHLORIDE; PSEUDOEPHEDRINE HYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL

CETIRIZINE HYDROCHLORIDE AND PSEUDOEPHEDRINE HYDROCHLORIDE

AUROBINDO PHARMA LTD	5MG;120MG	A212409	001	Mar 08, 2023
----------------------	-----------	---------	-----	--------------

CETRORELIX ACETATE

POWDER;SUBCUTANEOUS

CETROTIDE

+ EMD SERONO INC	EQ 3MG BASE/VIAL	N021197	002	Aug 11, 2000
------------------	------------------	---------	-----	--------------

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CETYL ALCOHOL; COLFOSCERIL PALMITATE; TYLOXAPOL

FOR SUSPENSION; INTRATRACHEAL

EXOSURF NEONATAL

GLAXOSMITHKLINE 12MG/VIAL; 108MG/VIAL; 8MG/VIAL N020044 001 Aug 02, 1990

CEVIMELINE HYDROCHLORIDE

CAPSULE; ORAL

CEVIMELINE HYDROCHLORIDE

APOTEX INC 30MG A091260 001 Aug 25, 2011

CHENODIOL

TABLET; ORAL

CHENIX

+ LEADIANT BIOSCI INC 250MG ** N018513 002 Jul 28, 1983

CHLOPHEDIANOL HYDROCHLORIDE

SYRUP; ORAL

ULO

3M 25MG/5ML N012126 001

CHLORAMPHENICOL

CREAM; TOPICAL

CHLOROMYCETIN

PARKE DAVIS 1% N050183 001

FOR SOLUTION; OPHTHALMIC

CHLOROMYCETIN

PARKEDALE 25MG/VIAL N050143 001

INJECTABLE; INJECTION

CHLOROMYCETIN

PARKE DAVIS 250MG/ML N050153 001

OINTMENT; OPHTHALMIC

CHLORAMPHENICOL

ALTANA 1% A060133 001

CHLOROFAIR

PHARMAFAIR 1% A062439 001 Apr 21, 1983

CHLOROMYCETIN

PARKEDALE 1% N050156 001

CHLOROPTIC S.O.P.

ALLERGAN 1% A061187 001

ECONOCHLOR

ALCON 1% A061648 001

SOLUTION/DROPS; OPHTHALMIC

CHLORAMPHENICOL

ALCON 0.5% A062628 001 Sep 25, 1985

EPIC PHARMA LLC 0.5% A062042 001

CHLOROFAIR

PHARMAFAIR 0.5% A062437 001 Apr 14, 1983

CHLOROPTIC

ALLERGAN 0.5% N050091 001

ECONOCHLOR

ALCON 0.5% A061645 001

OPHTHOCHLOR

PARKEDALE 0.5% A061220 001

OPTOMYCIN

OPTOPICS 0.5% A062171 001 Mar 31, 1982

SOLUTION/DROPS; OTIC

CHLOROMYCETIN

PARKEDALE 0.5% N050205 001

CHLORAMPHENICOL SODIUM SUCCINATE

INJECTABLE; INJECTION

CHLORAMPHENICOL

ELKINS SINN EQ 1GM BASE/VIAL A062406 001 Nov 09, 1982

CHLORAMPHENICOL SODIUM SUCCINATE

FRESENIUS KABI USA EQ 1GM BASE/VIAL A062365 001 Aug 25, 1982

GRUPPO LEPETIT EQ 1GM BASE/VIAL A062278 001

CHLOROMYCETIN

+ PARKEDALE EQ 1GM BASE/VIAL N050155 001

MYCHEL-S

ANGUS EQ 1GM BASE/VIAL A060132 001

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CHLORAMPHENICOL; HYDROCORTISONE ACETATE

FOR SUSPENSION;OPHTHALMIC

CHLOROMYCETIN HYDROCORTISONE

PARKEDALE	12.5MG/VIAL;25MG/VIAL	N050202	001
-----------	-----------------------	---------	-----

CHLORAMPHENICOL; HYDROCORTISONE ACETATE; POLYMYXIN B SULFATE

OINTMENT;OPHTHALMIC

OPHTHOCORT

PARKEDALE	10MG/GM;5MG/GM;10,000 UNITS/GM	N050201	002
-----------	--------------------------------	---------	-----

CHLORAMPHENICOL; POLYMYXIN B SULFATE

OINTMENT;OPHTHALMIC

CHLOROMYXIN

PARKE DAVIS	1%;10,000 UNITS/GM	N050203	002
-------------	--------------------	---------	-----

CHLORAMPHENICOL; PREDNISOLONE

OINTMENT;OPHTHALMIC

CHLOROPTIC-P S.O.P.

ALLERGAN	1%;0.5%	A061188	001
----------	---------	---------	-----

CHLORDIAZEPOXIDE

CAPSULE, EXTENDED RELEASE;ORAL

LIBRELEASE

VALEANT PHARM INTL	30MG	N017813	001	Sep 12, 1983
--------------------	------	---------	-----	--------------

TABLET;ORAL

LIBRITABS

VALEANT PHARM INTL	5MG	A085482	001
	10MG	A085481	001
	25MG	A085488	001

CHLORDIAZEPOXIDE HYDROCHLORIDE

CAPSULE;ORAL

A-POXIDE

ABBOTT	5MG	A085447	001
	5MG	A085517	001
	10MG	A085447	002
	10MG	A085518	001
	25MG	A085447	003
	25MG	A085513	001

CHLORDIAZACHEL

RACHELLE	5MG	A085086	001
	10MG	A084639	001
	25MG	A085087	001

CHLORDIAZEPOXIDE HYDROCHLORIDE

ASCOT	5MG	A087525	001	Jan 07, 1982
	10MG	A087524	001	Jan 07, 1982
	25MG	A087512	001	Jan 07, 1982
FERRANTE	5MG	A085118	001	
	10MG	A085119	001	
	25MG	A085120	001	
HALSEY	5MG	A085340	001	
	10MG	A085339	001	
	25MG	A084685	001	
IMPAX LABS	5MG	A086213	001	
	10MG	A085113	001	
	25MG	A086212	001	
IVAX SUB TEVA PHARMS	5MG	A083741	001	
	10MG	A083742	001	
	25MG	A083570	001	
LEDERLE	5MG	A086892	001	
	5MG	A087234	001	
	10MG	A086876	001	
	10MG	A087037	001	
	25MG	A086893	001	
	25MG	A087231	001	
MAST MM	10MG	A086217	001	
MYLAN	5MG	A084886	001	
	10MG	A084601	001	
	25MG	A084887	001	
PARKE DAVIS	5MG	A085163	001	
	10MG	A084598	001	
	25MG	A085164	001	

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CHLORDIAZEPOXIDE HYDROCHLORIDE

CAPSULE; ORAL

CHLORDIAZEPOXIDE HYDROCHLORIDE

PIONEER PHARMS	10MG	A089533 001	Jul 15, 1988
	25MG	A089558 001	Jul 15, 1988
PUREPAC PHARM	5MG	A085155 001	
	10MG	A084939 002	
	25MG	A085144 001	
ROXANE	5MG	A084706 001	
	10MG	A084700 001	
	25MG	A084705 001	
SUPERPHARM	5MG	A088987 001	Apr 25, 1985
	10MG	A088986 001	Apr 25, 1985
	25MG	A088988 001	Apr 25, 1985
TEVA	5MG	A088705 001	Jan 18, 1985
	10MG	A088706 001	Jan 18, 1985
	25MG	A086494 001	
	25MG	A088707 001	Jan 18, 1985
UPSHER SMITH LABS	5MG	A084919 001	
+	10MG	A084920 001	
	25MG	A084823 001	
USL PHARMA	5MG	A084644 001	
	10MG	A084623 001	
	25MG	A084645 001	
VANGARD	5MG	A088129 001	Mar 28, 1983
	10MG	A088010 001	Mar 28, 1983
	25MG	A088130 001	Mar 28, 1983
WATSON LABS	5MG	A086383 001	
	10MG	A086294 001	
	25MG	A086382 001	
WEST WARD	5MG	A085014 001	
	10MG	A085000 001	
	25MG	A085294 001	
LIBRIUM			
+	VALEANT PHARM INTL 5MG **	N012249 002	
+	10MG **	N012249 001	
+	25MG **	N012249 003	
LYGEN			
ALRA	5MG	A085107 001	
	10MG	A085009 001	
	25MG	A085108 001	

INJECTABLE; INJECTION

LIBRIUM

BAUSCH 100MG/AMP N012301 001

CHLORDIAZEPOXIDE HYDROCHLORIDE; CLIDINIUM BROMIDE

CAPSULE; ORAL

CHLORDIAZEPOXIDE HYDROCHLORIDE AND CLIDINIUM BROMIDE

TEVA PHARMS USA	5MG;2.5MG	A211476 001	Nov 02, 2021
TORRENT	5MG;2.5MG	A217385 001	Dec 14, 2023

CHLORDIAZEPOXIDE; ESTROGENS, ESTERIFIED

TABLET; ORAL

MENRIUM 10-4

ROCHE 10MG;0.4MG N014740 006

MENRIUM 5-2

ROCHE 5MG;0.2MG N014740 002

MENRIUM 5-4

ROCHE 5MG;0.4MG N014740 004

CHLORHEXIDINE GLUCONATE

SOLUTION; DENTAL

CHLORHEXIDINE GLUCONATE

SCIEGEN PHARMS INC 0.12% A074356 001 May 07, 1996

PAROEX

SUNSTAR AMERICAS 0.12% A076434 001 Nov 29, 2005

PERIOGARD

COLGATE PALMOLIVE CO 0.12% A073695 001 Jan 14, 1994

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CHLORHEXIDINE GLUCONATE

SOLUTION; TOPICAL

BRIAN CARE

SOAPCO

4%

A071419 001 Dec 17, 1987

EXIDINE

XTTRIUM

2.5%

N019421 001 Dec 17, 1985

MICRODERM

J AND J

4%

A072255 001 Apr 15, 1991

PREVACARE R

J AND J

0.5%

A072292 001 Jan 28, 1992

STERI-STAT

MATRIX MEDCL

4%

A070104 001 Jul 24, 1986

SPONGE; TOPICAL

CHLORHEXIDINE GLUCONATE

KENDALL IL

4%

N019490 001 Mar 27, 1987

E-Z SCRUB

BECTON DICKINSON

4%

A073416 001 Mar 14, 2000

HIBICLENS

+ MOLNLYCKE HLTH

4% **

N018423 001

MICRODERM

J AND J

4%

A072295 001 Feb 28, 1991

PHARMASEAL SCRUB CARE

CAREFUSION 2200

4%

N019793 001 Dec 02, 1988

TABLET; DENTAL

PERIOCHIP

+ DEXCEL PHARMA

2.5MG

N020774 001 May 15, 1998

CHLORMERODRIN HG-197

INJECTABLE; INJECTION

CHLORMERODRIN HG 197

BRACCO

0.6-1.4mCi/ML

N017269 001

CHLORMEZANONE

TABLET; ORAL

TRANCOPAL

SANOFI AVENTIS US

100MG

N011467 003

200MG

N011467 005

CHLOROPROCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

CHLOROPROCAINE HYDROCHLORIDE

HOSPIRA

2%

A087447 001 Apr 16, 1982

3%

A087446 001 Apr 16, 1982

NESACAINE

+ FRESENIUS KABI USA

1%

N009435 001

NESACAINE-MPF

FRESENIUS KABI USA

2%

N009435 003

3%

N009435 004

SOLUTION; INTRATHECAL

CLOROTEKAL

+ B BRAUN MEDICAL INC

50MG/5ML (10MG/ML)

N208791 001 Sep 26, 2017

CHLOROQUINE HYDROCHLORIDE

INJECTABLE; INJECTION

ARALEN HYDROCHLORIDE

SANOFI AVENTIS US

EQ 40MG BASE/ML

N006002 002

CHLOROQUINE PHOSPHATE

TABLET; ORAL

ARALEN

+ SANOFI AVENTIS US

500MG **

N006002 001

CHLOROQUINE PHOSPHATE

HIKMA PHARMS

250MG **

A083082 001

500MG **

A083082 002 Sep 17, 1999

IPCA LABS LTD

250MG

A090610 001 Dec 03, 2009

500MG

A090249 001 Dec 03, 2009

MD PHARM

250MG

A087228 001

PUREPAC PHARM

250MG

A080886 001

TEVA

250MG

A087504 001 Jan 13, 1982

WATSON LABS

250MG

A087979 001 Dec 21, 1982

500MG

A088030 001 Dec 21, 1982

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CHLOROQUINE PHOSPHATE; PRIMAQUINE PHOSPHATE

TABLET; ORAL

ARALEN PHOSPHATE W/ PRIMAQUINE PHOSPHATE

SANOFI AVENTIS US EQ 300MG BASE;EQ 45MG BASE N014860 002

CHLOROTHIAZIDE

TABLET; ORAL

CHLOROTHIAZIDE

ABC HOLDING	250MG	A085569	001	
HIKMA INTL PHARMS	250MG	A086028	001	Jul 14, 1982
	500MG	A087736	001	Jul 14, 1982
LEDERLE	250MG	A086940	001	
	500MG	A086938	001	
NORVIUM BIOSCIENCE	250MG	A084217	002	
	500MG	A084217	001	
SANDOZ	250MG	A085485	001	
WATSON LABS	250MG	A085165	001	
	250MG	A085173	001	
	250MG	A086795	001	Aug 15, 1983
	500MG	A084026	001	Sep 01, 1982
	500MG	A086796	001	Aug 15, 1983

DIURIL

+ RISING 250MG ** N011145 004

+ 500MG ** N011145 002

CHLOROTHIAZIDE SODIUM

INJECTABLE; INJECTION

CHLOROTHIAZIDE SODIUM

RK PHARMA	EQ 500MG BASE/VIAL	A202493	001	Jun 18, 2014
SUN PHARM	EQ 500MG BASE/VIAL	A091546	001	Jul 26, 2011

DIURIL

+ RISING EQ 500MG BASE/VIAL ** N011145 005

CHLOROTHIAZIDE; METHYLDOPA

TABLET; ORAL

ALDOCLOR-150

MERCK 150MG;250MG N016016 001

ALDOCLOR-250

MERCK 250MG;250MG N016016 002

METHYLDOPA AND CHLOROTHIAZIDE

PAR PHARM	150MG;250MG	A070783	001	Nov 06, 1987
	250MG;250MG	A070654	001	Nov 06, 1987

CHLOROTHIAZIDE; RESERPINE

TABLET; ORAL

CHLOROTHIAZIDE AND RESERPINE

HIKMA	250MG;0.125MG	A088557	001	Dec 22, 1983
	500MG;0.125MG	A088365	001	Dec 22, 1983

CHLOROTHIAZIDE W/ RESERPINE

WATSON LABS	250MG;0.125MG	A084853	001	
	500MG;0.125MG	A088151	001	Jun 09, 1983

CHLOROTHIAZIDE-RESERPINE

NORVIUM BIOSCIENCE	250MG;0.125MG	A087744	001	May 06, 1982
	500MG;0.125MG	A087745	001	May 06, 1982

DIUPRES-250

MERCK 250MG;0.125MG N011635 003 Aug 26, 1987

DIUPRES-500

MERCK 500MG;0.125MG N011635 006 Aug 26, 1987

CHLOROTRIANISENE

CAPSULE; ORAL

CHLOROTRIANISENE

BANNER PHARMACAPS 12MG A084652 001

TACE

SANOFI AVENTIS US	12MG	N008102	004	
	25MG	N011444	001	
	72MG	N016235	001	

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CHLOROXYLINE

SHAMPOO; TOPICAL

CAPITROL

WESTWOOD SQUIBB

2%

N017594 001

CHLORPHENESIN CARBAMATE

TABLET; ORAL

MAOLATE

PAMLAB LLC

400MG

N014217 002

CHLORPHENIRAMINE MALEATE

CAPSULE, EXTENDED RELEASE; ORAL

CHLORPHENIRAMINE MALEATE

AUROLIFE PHARMA LLC

12MG

A070797 001 Aug 12, 1988

TELDRIN

GLAXOSMITHKLINE

8MG

N017369 001

12MG

N017369 002

INJECTABLE; INJECTION

CHLOR-TRIMETON

SCHERING PLOUGH

10MG/ML

N008826 001

100MG/ML

N008794 001

CHLORPHENIRAMINE MALEATE

BEL MAR

10MG/ML

A080821 001

ELKINS SINN

10MG/ML

A080797 001

WATSON LABS

10MG/ML

A083593 001

10MG/ML

A086096 001

100MG/ML

A086095 001

PYRIDAMAL 100

BEL MAR

100MG/ML

A083733 001

SYRUP; ORAL

CHLOR-TRIMETON

SCHERING

2MG/5ML

N006921 006

CHLORPHENIRAMINE MALEATE

PHARM ASSOC

2MG/5ML

A087520 001 Feb 10, 1982

TABLET; ORAL

ANTAGONATE

BAYER PHARMS

4MG

A083381 001

CHLOR-TRIMETON

SCHERING

4MG

N006921 002

CHLORPHENIRAMINE MALEATE

ANABOLIC

4MG

A083078 001

AUROLIFE PHARMA LLC

4MG

A080961 001

BELL PHARMA

4MG

A083062 001

ELKINS SINN

4MG

A080938 001

IMPAX LABS

4MG

A080809 001

IVAX SUB TEVA PHARMS

4MG

A080779 001

KV PHARM

4MG

A087164 001

LEDERLE

4MG

A086941 001

NEWTRON PHARMS

4MG

A086519 001

PANRAY

4MG

A083243 001

PHARMAVITE

4MG

A085104 001

PHARMERAL

4MG

A083753 001

PIONEER PHARMS

4MG

A088556 001 Jul 13, 1984

PUREPAC PHARM

4MG

A086306 001

PVT FORM

4MG

A080786 001

ROXANE

4MG

A080626 001

SUN PHARM INDUSTRIES

4MG

A080700 001

VITARINE

4MG

A085837 001

WATSON LABS

4MG

A080696 001

4MG

A080791 001

4MG

A085139 001

WEST WARD

4MG

A083787 001

KLOROMIN

HALSEY

4MG

A083629 001

PHENETRON

LANNETT

4MG

A080846 001

TABLET, EXTENDED RELEASE; ORAL

CHLOR-TRIMETON

+ BAYER HEALTHCARE LLC

8MG **

N007638 001

+

12MG **

N007638 002

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CHLORPHENIRAMINE MALEATE

TABLET, EXTENDED RELEASE;ORAL

EFIDAC 24 CHLORPHENIRAMINE MALEATE

ALZA

16MG

N019746 002 Nov 18, 1994

CHLORPHENIRAMINE MALEATE; HYDROCODONE BITARTRATE

SOLUTION;ORAL

HYDROCODONE BITARTRATE AND CHLORPHENIRAMINE MALEATE

ACELLA

4MG/5ML;5MG/5ML

A205691 001 Jun 09, 2017

TRIS PHARMA INC

4MG/5ML;5MG/5ML

A206438 001 Jan 27, 2015

VITUZ

+ PERSION

4MG/5ML;5MG/5ML

N204307 001 Feb 20, 2013

CHLORPHENIRAMINE MALEATE; HYDROCODONE BITARTRATE; PSEUDOEPHEDRINE HYDROCHLORIDE

SOLUTION;ORAL

HYDROCODONE BITARTRATE, CHLORPHENIRAMINE MALEATE AND PSEUDOEPHEDRINE HYDROCHLORIDE

MAYNE PHARMA INC

4MG/5ML;5MG/5ML;60MG/5ML

A205657 001 Aug 03, 2015

SCIEGEN PHARMS INC

4MG/5ML;5MG/5ML;60MG/5ML

A206660 001 May 15, 2017

TRIS PHARMA INC

4MG/5ML;5MG/5ML;60MG/5ML

A203838 001 Nov 26, 2014

ZUTRIPRO

+ PERSION

4MG/5ML;5MG/5ML;60MG/5ML **

N022439 001 Jun 08, 2011

CHLORPHENIRAMINE MALEATE; PHENYLPROPANOLAMINE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE;ORAL

COLD CAPSULE IV

GRAHAM DM

12MG;75MG

N018793 001 Apr 25, 1985

COLD CAPSULE V

GRAHAM DM

8MG;75MG

N018794 001 Apr 23, 1985

TABLET, EXTENDED RELEASE;ORAL

TRIAMINIC-12

NOVARTIS

12MG;75MG

N018115 001

CHLORPHENIRAMINE MALEATE; PSEUDOEPHEDRINE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE;ORAL

CODIMAL-L.A. 12

SCHWARZ PHARMA

12MG;120MG

N018935 001 Apr 15, 1985

ISOCOLOR

FISONS

8MG;120MG

N018747 001 Mar 06, 1986

PSEUDOEPHEDRINE HYDROCHLORIDE AND CHLORPHENIRAMINE MALEATE

CENT PHARMS

8MG;120MG

N019428 001 Aug 02, 1988

GRAHAM DM

8MG;120MG

N018844 001 Mar 20, 1985

12MG;120MG

N018843 001 Mar 18, 1985

KV PHARM

12MG;120MG

A071455 001 Mar 01, 1989

CHLORPHENIRAMINE MALEATE; PSEUDOEPHEDRINE SULFATE

TABLET, EXTENDED RELEASE;ORAL

CHLOR-TRIMETON

+ BAYER HEALTHCARE LLC

8MG;120MG

N018397 001

CHLORPHENIRAMINE POLISTIREX; CODEINE POLISTIREX

SUSPENSION, EXTENDED RELEASE;ORAL

CODEPREX

LANNETT CO INC

EQ 4MG MALEATE/5ML;EQ 20MG BASE/5ML

N021369 001 Jun 21, 2004

PENNTUSS

FISONS

EQ 4MG MALEATE/5ML;EQ 10MG BASE/5ML

N018928 001 Aug 14, 1985

TUZISTRA XR

+ TRIS PHARMA INC

EQ 2.8MG BASE/5ML;EQ 14.7MG BASE/5ML

N207768 001 Apr 30, 2015

CHLORPHENIRAMINE POLISTIREX; HYDROCODONE POLISTIREX

CAPSULE, EXTENDED RELEASE;ORAL

TUSSICAPS

ECR PHARMA

EQ 4MG MALEATE;EQ 5MG BITARTRATE

A077273 002 Sep 24, 2007

EQ 8MG MALEATE;EQ 10MG BITARTRATE

A077273 001 Sep 24, 2007

SUSPENSION, EXTENDED RELEASE;ORAL

HYDROCODONE POLISTIREX AND CHLORPHENIRAMINE POLISTIREX

NEOS THERAPS INC

EQ 8MG MALEATE/5ML;EQ 10MG

A091671 001 Jun 29, 2012

BITARTRATE/5ML

TUSSIONEX PENNKINETIC

+ UCB INC

EQ 8MG MALEATE/5ML;EQ 10MG

N019111 001 Dec 31, 1987

BITARTRATE/5ML **

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CHLORPHENTERMINE HYDROCHLORIDE

TABLET; ORAL

PRE-SATE

PARKE DAVIS EQ 65MG BASE N014696 001

CHLORPROMAZINE

SUPPOSITORY; RECTAL

THORAZINE

+ GLAXOSMITHKLINE 25MG ** N009149 024

+ 100MG ** N009149 033

CHLORPROMAZINE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

THORAZINE

GLAXOSMITHKLINE 30MG N011120 016

75MG N011120 017

150MG N011120 018

200MG N011120 019

300MG N011120 020

CONCENTRATE; ORAL

CHLORPROMAZINE HYDROCHLORIDE

ACTAVIS MID ATLANTIC 100MG/ML A086863 001

PHARM ASSOC 30MG/ML A040231 001 Dec 30, 1999

100MG/ML A040224 001 Jan 26, 1999

WOCKHARDT 30MG/ML A087032 001 Jul 08, 1982

100MG/ML A087053 001

CHLORPROMAZINE HYDROCHLORIDE INTENSOL

HIKMA 30MG/ML A088157 001 Apr 27, 1983

100MG/ML A088158 001 Apr 27, 1983

SONAZINE

FOSUN PHARMA 30MG/ML A080983 004

100MG/ML A080983 005

THORAZINE

+ GLAXOSMITHKLINE 30MG/ML ** N009149 032

+ 100MG/ML ** N009149 043

INJECTABLE; INJECTION

CHLORPROMAZINE HYDROCHLORIDE

ABRAXIS PHARM 25MG/ML A084911 001

MARSAM PHARMS LLC 25MG/ML A089563 001 Apr 15, 1988

WATSON LABS 25MG/ML A085591 001

WYETH AYERST 25MG/ML A080370 001

THORAZINE

+ GLAXOSMITHKLINE 25MG/ML ** N009149 011

SYRUP; ORAL

CHLORPROMAZINE HYDROCHLORIDE

ALPHARMA US PHARMS 10MG/5ML A086712 001

SONAZINE

CHARTWELL RX 10MG/5ML A083040 001

THORAZINE

+ GLAXOSMITHKLINE 10MG/5ML ** N009149 022

TABLET; ORAL

CHLORPROMAZINE HYDROCHLORIDE

ABBOTT 10MG A084414 001

25MG A084415 001

50MG A084411 001

100MG A084412 001

200MG A084413 001

CHARTWELL RX 10MG ** A080439 001

25MG ** A080439 002

50MG ** A080439 003

100MG ** A080439 004

200MG ** A080439 005

CYCLE 10MG A085331 001

25MG A085331 002

50MG A085331 003

100MG A085331 004

200MG A085331 005

IVAX SUB TEVA PHARMS 10MG A083549 001

25MG A083549 002

50MG A083549 003

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CHLORPROMAZINE HYDROCHLORIDE

TABLET;ORAL

CHLORPROMAZINE HYDROCHLORIDE

	100MG	A083574	001	
	200MG	A083575	001	
KV PHARM	10MG	A085750	002	Jan 04, 1982
	25MG	A085751	001	
	50MG	A085484	001	
	100MG	A085752	001	
	200MG	A085748	002	Jan 04, 1982
LEDERLE	10MG	A084803	001	
	25MG	A084801	001	
	50MG	A084800	001	
	100MG	A084789	001	
	200MG	A084802	001	
PUREPAC PHARM	10MG	A080403	004	
	25MG	A080403	001	
	50MG	A080403	002	
	100MG	A080403	003	
	200MG	A080403	005	
PVT FORM	25MG	A080340	001	
	50MG	A080340	002	
	200MG	A080340	003	
VANGARD	10MG	A088038	001	Aug 16, 1982
	25MG	A087645	001	
	50MG	A087646	001	
WATSON LABS	10MG	A085959	001	
	25MG	A085956	001	
	50MG	A085960	001	
	100MG	A085957	001	
	200MG	A085958	001	
WEST WARD	10MG	A087783	001	Sep 16, 1982
	25MG	A087865	001	Sep 16, 1982
	50MG	A087878	001	Sep 15, 1982
	100MG	A087884	001	Sep 15, 1982
	200MG	A087880	001	Sep 16, 1982
PROMAPAR				
PARKE DAVIS	10MG	A086886	001	
	25MG	A084423	001	
	50MG	A086887	001	
	100MG	A086888	001	
	200MG	A086885	001	
THORAZINE				
GLAXOSMITHKLINE	10MG **	N009149	002	
	25MG **	N009149	007	
	50MG **	N009149	013	
	100MG **	N009149	018	
	200MG **	N009149	020	

CHLORPROPAMIDE

TABLET;ORAL

CHLORPROPAMIDE

ANI PHARMS	100MG	A088768	001	Oct 11, 1984
	100MG	A088812	001	Oct 19, 1984
	100MG	A088840	001	Oct 25, 1984
	100MG	A088918	001	Oct 16, 1984
	100MG	A088921	001	Apr 12, 1985
	100MG	A089446	001	Nov 17, 1986
	250MG	A087353	001	
	250MG	A088813	001	Oct 19, 1984
	250MG	A088826	001	Sep 26, 1984
	250MG	A088919	001	Oct 16, 1984
	250MG	A088922	001	Apr 12, 1985
	250MG	A089447	001	Nov 17, 1986
DAVA PHARMS INC	100MG	A089561	001	Sep 04, 1987
	250MG	A089562	001	Sep 04, 1987
HALSEY	100MG	A089321	001	Jan 16, 1986
	250MG	A088662	001	Jan 09, 1986
NORVIUM BIOSCIENCE	100MG	A088549	002	Jun 01, 1984
	250MG	A088549	001	Jun 01, 1984

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CHLORPROPAMIDE

TABLET; ORAL

CHLORPROPAMIDE

PAR PHARM	100MG	A088175 001	Feb 27, 1984
	250MG	A088176 001	Feb 27, 1984
RISING	100MG	A088725 001	Aug 31, 1984
	250MG	A088726 001	Aug 31, 1984
SANDOZ	250MG	A084669 001	
SUPERPHARM	100MG	A088694 001	Sep 17, 1984
	250MG	A088695 001	Sep 17, 1984
USL PHARMA	100MG	A088708 001	Aug 30, 1984
	250MG	A088709 001	Aug 30, 1984
WATSON LABS	100MG	A086865 001	Sep 24, 1984
	100MG	A088608 001	Apr 12, 1984
	250MG	A086866 001	
	250MG	A088568 001	Apr 12, 1984
WATSON LABS TEVA	100MG	A088852 001	Sep 26, 1984
DIABINESE			
+ PFIZER	100MG	N011641 003	
+	250MG	N011641 006	
GLUCAMIDE			
ANI PHARMS	250MG	A088641 001	Oct 11, 1984

CHLORPROTHIXENE

CONCENTRATE; ORAL

TARACTAN

ROCHE	100MG/5ML	N016149 002	
INJECTABLE; INJECTION			
TARACTAN			
ROCHE	12.5MG/ML	N012487 001	
TABLET; ORAL			
TARACTAN			
ROCHE	10MG	N012486 005	
	25MG	N012486 004	
	50MG	N012486 003	
	100MG	N012486 001	

CHLORTETRACYCLINE HYDROCHLORIDE

OINTMENT; OPHTHALMIC

AUREOMYCIN

LEDERLE	1%	N050404 001	
---------	----	-------------	--

CHLORTHALIDONE

TABLET; ORAL

CHLORTHALIDONE

ABBOTT	25MG	A087364 001	
	50MG	A087384 001	
ANI PHARMS	25MG	A087296 001	
	25MG	A087706 001	
	25MG	A088164 001	Jan 09, 1984
	50MG	A087689 001	
ASCOT	25MG	A087698 001	Oct 20, 1982
	50MG	A087699 001	Oct 20, 1982
BARR LABS INC	25MG	A088902 001	Sep 19, 1985
	50MG	A088903 001	Sep 19, 1985
COSETTE	50MG	A088651 001	May 30, 1985
DAVA PHARMS INC	25MG	A087451 001	
	50MG	A087450 001	
IVAX PHARMS	25MG	A087555 001	
	50MG	A087176 001	
	50MG	A087947 001	Feb 27, 1984
KV PHARM	25MG	A087311 001	
	50MG	A087312 001	
MUTUAL PHARM	25MG	A087292 001	
	25MG	A089738 001	Sep 19, 1988
	50MG	A087293 001	
	50MG	A089739 001	Sep 19, 1988
PIONEER PHARMS	50MG	A089591 001	Jul 21, 1988
PUREPAC PHARM	25MG	A088139 001	Jul 16, 1986
	50MG	A088140 001	Aug 11, 1983
SANDOZ	25MG	A087380 001	

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CHLORTHALIDONE

TABLET; ORAL

CHLORTHALIDONE

	50MG	A087118	001	
	50MG	A087381	001	
SUNNY	25MG	A209068	001	Jan 25, 2022
	50MG	A209068	002	Jan 25, 2022
SUPERPHARM	25MG	A087473	001	Feb 09, 1983
	50MG	A087247	001	Feb 09, 1983
USL PHARMA	25MG	A089051	001	Jun 01, 1987
	50MG	A089052	001	Jun 01, 1987
VANGARD	25MG	A088012	001	Jul 14, 1982
	50MG	A088073	001	Mar 25, 1983
WARNER CHILCOTT	25MG	A087515	001	Jan 24, 1983
	50MG	A087516	001	Feb 09, 1983
WATSON LABS	25MG	A087050	001	
	25MG	A087100	001	
	50MG	A087029	001	
	50MG	A087082	001	
	50MG	A087521	001	
HYGROTON				
+ SANOFI AVENTIS US	25MG **	N012283	004	
+	50MG **	N012283	003	
THALITONE				
MONARCH PHARMS	25MG	A088051	001	Nov 12, 1982

CHLORTHALIDONE; CLONIDINE HYDROCHLORIDE

TABLET; ORAL

CLONIDINE HYDROCHLORIDE AND CHLORTHALIDONE

PAR PHARM	15MG; 0.1MG	A071179	001	Dec 16, 1987
	15MG; 0.2MG	A071178	001	Dec 16, 1987
	15MG; 0.3MG	A071142	001	Dec 16, 1987
CLORPRES				
NATCO PHARMA	15MG; 0.1MG	A071325	003	Feb 09, 1987
	15MG; 0.2MG	A071325	002	Feb 09, 1987
	15MG; 0.3MG	A071325	001	Feb 09, 1987
COMBIPRES				
+ BOEHRINGER INGELHEIM	15MG; 0.1MG **	N017503	001	
+	15MG; 0.2MG **	N017503	002	
+	15MG; 0.3MG **	N017503	003	Apr 10, 1984

CHLORTHALIDONE; METOPROLOL TARTRATE

CAPSULE; ORAL

LOPRESSIDONE

NOVARTIS	25MG; 100MG	N019451	001	Dec 31, 1987
	25MG; 200MG	N019451	002	Dec 31, 1987

CHLORTHALIDONE; RESERPINE

TABLET; ORAL

DEMI-REGROTON

SANOFI AVENTIS US	25MG; 0.125MG	N015103	002	
REGROTON				
SANOFI AVENTIS US	50MG; 0.25MG	N015103	001	

CHLORZOXAZONE

TABLET; ORAL

CHLORZOXAZONE

ACME LABS	500MG	A214365	001	Apr 27, 2023
ACTAVIS ELIZABETH	250MG	A088928	001	May 08, 1987
	500MG	A040113	001	Sep 29, 1995
BARR	500MG	A089895	001	May 04, 1988
CHARTWELL RX	375MG	A212053	001	Sep 14, 2020
	750MG	A212053	002	Sep 14, 2020
GLENMARK PHARMS LTD	375MG	A212185	001	May 26, 2020
	750MG	A212185	002	May 26, 2020
OHM LABS	250MG	A081298	001	Dec 29, 1993
	500MG	A081299	001	Dec 29, 1993
PIONEER PHARMS	250MG	A089592	001	Jan 06, 1989
	500MG	A089948	001	Jan 06, 1989
RISING	250MG	A089852	001	May 04, 1988
STRIDES PHARMA	250MG	A087981	001	Sep 20, 1983

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CHLORZOXAZONE

TABLET; ORAL

CHLORZOXAZONE

SUN PHARM INDUSTRIES	500MG	A089970	001	Sep 27, 1990
TEVA PHARMS USA INC	375MG	A212898	001	Jun 17, 2020
	750MG	A212898	002	Jun 17, 2020
WATSON LABS	250MG	A086901	001	
	250MG	A086948	001	Aug 09, 1982
	500MG	A040137	001	Aug 09, 1996
	500MG	A081019	001	Jul 29, 1991
	500MG	A081040	001	Aug 22, 1989
PARAFLEX				
+ ORTHO MCNEIL PHARM	250MG **	N011300	003	
PARAFON FORTE DSC				
+ JANSSEN R AND D	500MG **	N011529	002	Jun 15, 1987
STRIFON FORTE DSC				
FERNDALE LABS	500MG	A081008	001	Dec 23, 1988

CHOLESTYRAMINE

BAR, CHEWABLE; ORAL

CHOLYBAR

PARKE DAVIS	EQ 4GM RESIN/BAR	A071621	001	May 26, 1988
	EQ 4GM RESIN/BAR	A071739	001	May 26, 1988

POWDER; ORAL

CHOLESTYRAMINE

ANI PHARMS	EQ 4GM RESIN/PACKET	A074554	001	Oct 02, 1996
	EQ 4GM RESIN/SCOOPFUL	A074554	002	Oct 02, 1996
IVAX SUB TEVA PHARMS	EQ 4GM RESIN/PACKET	A074771	001	Jul 09, 1997
	EQ 4GM RESIN/SCOOPFUL	A074771	002	Jul 09, 1997
TEVA	EQ 4GM RESIN/PACKET	A074347	001	May 28, 1998
	EQ 4GM RESIN/SCOOPFUL	A074347	002	May 28, 1998
UPSHER SMITH LABS	EQ 4GM RESIN/PACKET	A214877	001	Jan 21, 2022
	EQ 4GM RESIN/SCOOPFUL	A214877	002	Jan 21, 2022

CHOLESTYRAMINE LIGHT

TEVA	EQ 4GM RESIN/PACKET	A074348	001	May 28, 1998
	EQ 4GM RESIN/SCOOPFUL	A074348	002	May 28, 1998
TEVA PHARMS	EQ 4GM RESIN/PACKET	A074555	001	Sep 30, 1998
	EQ 4GM RESIN/SCOOPFUL	A074555	002	Sep 30, 1998

QUESTRAN

+ BRISTOL MYERS	EQ 4GM RESIN/PACKET **	N016640	001	
+	EQ 4GM RESIN/SCOOPFUL **	N016640	003	

QUESTRAN LIGHT

+ BRISTOL MYERS	EQ 4GM RESIN/PACKET **	N019669	001	Dec 05, 1988
+	EQ 4GM RESIN/SCOOPFUL **	N019669	003	Dec 05, 1988

TABLET; ORAL

QUESTRAN

APOTHECON	EQ 800MG RESIN	A073403	002	Dec 27, 1999
	EQ 1GM RESIN	A073403	001	Apr 28, 1994

CHOLINE C-11

INJECTABLE; INTRAVENOUS

CHOLINE C-11

UCSF RODIOPHARM	4-33.1mCi/ML	A208444	001	Nov 20, 2017
UNIV TX MD ANDERSON	4-100mCi/ML	A205690	001	Oct 29, 2015
WA UNIV SCH MED	4-33.1mCi/ML	A208413	001	Jan 10, 2017

CHOLINE FENOFIBRATE

CAPSULE, DELAYED RELEASE; ORAL

FENOFIBRIC ACID

ENDO OPERATIONS	EQ 45MG FENOFIBRIC ACID	A201573	002	Jul 18, 2013
	EQ 135MG FENOFIBRIC ACID	A201573	001	Jul 18, 2013
MACLEODS PHARMS LTD	EQ 45MG FENOFIBRIC ACID	A207796	001	Feb 08, 2024
	EQ 135MG FENOFIBRIC ACID	A207796	002	Feb 08, 2024
NORVIUM BIOSCIENCE	EQ 45MG FENOFIBRIC ACID	A200913	001	Mar 25, 2013
	EQ 135MG FENOFIBRIC ACID	A200913	002	Mar 25, 2013
TWI PHARMS	EQ 45MG FENOFIBRIC ACID	A210469	001	Jul 05, 2019
	EQ 135MG FENOFIBRIC ACID	A210469	002	Jul 05, 2019

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CHROMIC CHLORIDE

INJECTABLE; INJECTION

CHROMIC CHLORIDE

ABRAXIS PHARM

EQ 0.004MG CHROMIUM/ML

N019271 001 May 05, 1987

CHROMIC PHOSPHATE P-32

INJECTABLE; INJECTION

PHOSPHOCOL P32

CURIUM

5mCi/ML

N017084 001

CICLESONIDE

AEROSOL, METERED; NASAL

ZETONNA

+ COVIS

0.037MG/INH

N202129 001 Jan 20, 2012

CICLOPIROX

GEL; TOPICAL

CICLOPIROX

+ ALVOGEN

0.77% **

N020519 001 Jul 21, 1997

SHAMPOO; TOPICAL

CICLOPIROX

ENCUBE

1%

A209975 001 Apr 05, 2018

SOLUTION; TOPICAL

CICLOPIROX

COSETTE

8%

A078233 001 Sep 18, 2007

EPIC PHARMA LLC

8%

A078975 001 Feb 17, 2010

HIKMA

8%

A078270 001 Sep 18, 2007

MYLAN PHARMS INC

8%

A078567 001 Sep 18, 2007

TEVA PHARMS

8%

A078079 001 Sep 18, 2007

PENLAC

+ VALEANT BERMUDA

8% **

N021022 001 Dec 17, 1999

CIDOFOVIR

INJECTABLE; INJECTION

VISTIDE

+ GILEAD SCIENCES INC

EQ 75MG BASE/ML **

N020638 001 Jun 26, 1996

CILASTATIN SODIUM; IMIPENEM

INJECTABLE; INJECTION

PRIMAXIN

MERCK

EQ 250MG BASE/VIAL; 250MG/VIAL

A062756 001 Jan 08, 1987

EQ 500MG BASE/VIAL; 500MG/VIAL

A062756 002 Jan 08, 1987

POWDER; INTRAMUSCULAR

PRIMAXIN

MERCK

EQ 500MG BASE/VIAL; 500MG/VIAL

N050630 001 Dec 14, 1990

EQ 750MG BASE/VIAL; 750MG/VIAL

N050630 002 Dec 14, 1990

POWDER; INTRAVENOUS

IMIPENEM AND CILASTATIN

HOSPIRA INC

EQ 250MG BASE/VIAL; 250MG/VIAL

A090825 001 Nov 16, 2011

EQ 500MG BASE/VIAL; 500MG/VIAL

A090825 002 Nov 16, 2011

EQ 500MG BASE/VIAL; 500MG/VIAL

A091007 001 Nov 16, 2011

PRIMAXIN

+ MERCK

EQ 250MG BASE/VIAL; 250MG/VIAL **

N050587 001 Nov 26, 1985

CILOSTAZOL

TABLET; ORAL

CILOSTAZOL

ACTAVIS ELIZABETH

100MG

A077028 002 Nov 26, 2004

AUROBINDO PHARMA USA

50MG

A077019 001 Nov 23, 2004

100MG

A077019 002 Nov 23, 2004

CHARTWELL RX

50MG

A077310 001 Nov 08, 2005

100MG

A077021 001 Nov 23, 2004

HIKMA

50MG

A077024 001 May 17, 2005

100MG

A077024 002 May 17, 2005

IVAX SUB TEVA PHARMS

100MG

A077020 002 Mar 01, 2005

NOSTRUM LABS INC

50MG

A077708 001 Sep 28, 2009

100MG

A077708 002 Sep 28, 2009

PLIVA HRVATSKA DOO

50MG

A077898 001 Oct 29, 2007

100MG

A077898 002 Oct 29, 2007

RISING

50MG

A077323 002 Apr 20, 2006

100MG

A077323 001 Apr 20, 2006

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CILOSTAZOL

TABLET; ORAL

PLETAL

+	OTSUKA	50MG **	N020863 001	Jan 15, 1999
+		100MG **	N020863 002	Jan 15, 1999

CIMETIDINE

SUSPENSION; ORAL

TAGAMET HB 200

	GLAXOSMITHKLINE	200MG/20ML	N020951 001	Jul 09, 1999
--	-----------------	------------	-------------	--------------

TABLET; ORAL

CIMETIDINE

	CHARTWELL RX	200MG	A074100 001	Jan 31, 1995
		300MG	A074100 002	Jan 31, 1995
		400MG	A074100 003	Jan 31, 1995
		800MG	A074100 004	Jan 31, 1995
	CONTRACT PHARMACAL	200MG	A074961 001	Jun 19, 1998
		200MG	A074963 001	Jun 19, 1998
	CYCLE	300MG	A074361 001	Dec 23, 1994
		400MG	A074361 002	Dec 23, 1994
		800MG	A074371 001	Dec 23, 1994
	HIKMA	200MG	A074890 001	Dec 18, 1998
		300MG	A074890 002	Dec 18, 1998
		400MG	A074890 003	Dec 18, 1998
		800MG	A074890 004	Dec 18, 1998
	IVAX SUB TEVA PHARMS	200MG	A074401 001	May 30, 1995
		200MG	A074424 001	Jul 28, 1995
		200MG	A075345 001	Jun 16, 1999
		300MG	A074401 002	May 30, 1995
		300MG	A074424 002	Jul 28, 1995
		400MG	A074401 003	May 30, 1995
		400MG	A074424 003	Jul 28, 1995
		800MG	A074402 001	May 30, 1995
		800MG	A074424 004	Jul 28, 1995
	NOVITIUM PHARMA	300MG	A074340 001	Jun 23, 1995
		400MG	A074340 002	Jun 23, 1995
		800MG	A074339 001	Jun 23, 1995
	PERRIGO	100MG	A074972 001	Jun 19, 1998
	PLIVA	200MG	A074568 001	Feb 27, 1997
		300MG	A074568 002	Feb 27, 1997
		400MG	A074568 003	Feb 27, 1997
		800MG	A074566 001	Feb 27, 1997
	SANDOZ	100MG	A075122 001	Jun 19, 1998
		200MG	A074250 001	Jun 29, 1995
		200MG	A075122 002	Jun 19, 1998
		300MG	A074250 002	Jun 29, 1995
		400MG	A074250 003	Jun 29, 1995
		800MG	A074250 004	Jun 29, 1995
	TEVA	200MG	A074365 001	Feb 28, 1995
		300MG	A074365 002	Feb 28, 1995
		400MG	A074365 003	Feb 28, 1995
		800MG	A074365 004	Feb 28, 1995
	UPSHER SMITH LABS	200MG	A074506 001	Jan 24, 1996
		300MG	A074506 002	Jan 24, 1996
		400MG	A074506 003	Jan 24, 1996
		800MG	A074506 004	Jan 24, 1996
	WATSON LABS INC	200MG	A074349 001	Aug 30, 1996
		300MG	A074349 002	Aug 30, 1996
		400MG	A074349 003	Aug 30, 1996
		800MG	A074316 001	Feb 28, 1996
	WATSON LABS TEVA	200MG	A075425 001	Jul 29, 1999
	TAGAMET			
+	GLAXOSMITHKLINE	200MG **	N017920 002	
+		300MG **	N017920 003	
+		400MG **	N017920 004	Dec 14, 1983
+		800MG **	N017920 005	Apr 30, 1986
	TAGAMET HB			
+	MEDTECH PRODUCTS	100MG **	N020238 001	Jun 19, 1995

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CIMETIDINE HYDROCHLORIDE

INJECTABLE; INJECTION

CIMETIDINE HYDROCHLORIDE

COSETTE	EQ 300MG BASE/2ML	A074296 001	Mar 28, 1997
ENDO OPERATIONS	EQ 300MG BASE/2ML	A074005 001	Aug 31, 1994
	EQ 300MG BASE/2ML	A074428 001	Apr 25, 1996
HOSPIRA	EQ 300MG BASE/2ML	A074344 001	Jan 31, 1995
	EQ 300MG BASE/2ML	A074345 001	Jan 31, 1995
	EQ 300MG BASE/2ML	A074412 001	Mar 28, 1997
	EQ 300MG BASE/2ML	A074422 001	Jan 31, 1995
LUITPOLD	EQ 300MG BASE/2ML	A074353 001	Dec 20, 1994
TEVA PARENTERAL	EQ 300MG BASE/2ML	A074252 001	Nov 26, 1997
CIMETIDINE HYDROCHLORIDE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER			
HOSPIRA	EQ 6MG BASE/ML	A074269 001	Dec 27, 1994
	EQ 90MG BASE/100ML	A074468 005	Dec 29, 1994
	EQ 120MG BASE/100ML	A074468 006	Dec 29, 1994
	EQ 180MG BASE/100ML	A074468 003	Dec 29, 1994
	EQ 240MG BASE/100ML	A074468 004	Dec 29, 1994
	EQ 360MG BASE/100ML	A074468 001	Dec 29, 1994
	EQ 480MG BASE/100ML	A074468 002	Dec 29, 1994
TAGAMET			
GLAXOSMITHKLINE	EQ 300MG BASE/2ML **	N017939 002	
TAGAMET HYDROCHLORIDE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER			
+ GLAXOSMITHKLINE	EQ 6MG BASE/ML **	N019434 001	Oct 31, 1985
SOLUTION; ORAL			
CIMETIDINE HYDROCHLORIDE			
ANI PHARMS	EQ 300MG BASE/5ML	A074610 001	Sep 26, 1996
	EQ 300MG BASE/5ML	A074859 001	Jul 09, 1998
	EQ 300MG BASE/5ML	A075110 001	Jun 18, 1998
CYCLE	EQ 300MG BASE/5ML	A074541 001	Aug 05, 1997
G AND W LABS INC	EQ 300MG BASE/5ML	A074176 001	Jun 01, 1994
PHARM ASSOC	EQ 300MG BASE/5ML	A074553 001	Jan 27, 1997
	EQ 300MG BASE/5ML	A075560 001	Mar 15, 2000
PHARMOBEDIENT CNSLTG	EQ 300MG BASE/5ML	A074757 001	Oct 17, 1997
TAGAMET			
GLAXOSMITHKLINE	EQ 300MG BASE/5ML **	N017924 001	

CINACALCET HYDROCHLORIDE

TABLET; ORAL

CINACALCET HYDROCHLORIDE

LUPIN LTD	EQ 30MG BASE	A210548 001	Jun 28, 2019
	EQ 60MG BASE	A210548 002	Jun 28, 2019
	EQ 90MG BASE	A210548 003	Jun 28, 2019
NORVIUM BIOSCIENCE	EQ 30MG BASE	A203422 001	Oct 16, 2018
	EQ 60MG BASE	A203422 002	Oct 16, 2018
	EQ 90MG BASE	A203422 003	Oct 16, 2018
STEVENS J	30MG	A204364 001	Dec 02, 2021
	60MG	A204364 002	Dec 02, 2021
	90MG	A204364 003	Dec 02, 2021
SUN PHARM	EQ 30MG BASE	A207008 001	Oct 11, 2018
	EQ 90MG BASE	A207008 003	Oct 11, 2018
SENSIPAR			
+ AMGEN	EQ 30MG BASE	N021688 001	Mar 08, 2004
+	EQ 60MG BASE	N021688 002	Mar 08, 2004
+	EQ 90MG BASE	N021688 003	Mar 08, 2004

CINOXACIN

CAPSULE; ORAL

CINOXACIN

LILLY	250MG	N018067 001	
	500MG	N018067 002	
CINOXACIN			
TEVA	250MG	A073005 001	Feb 28, 1992
	500MG	A073006 001	Feb 28, 1992

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CIPROFLOXACIN

INJECTABLE; INJECTION

CIPRO

+	BAYER HLTHCARE	400MG/40ML (10MG/ML) **	N019847	001	Dec 26, 1990
+		200MG/20ML (10MG/ML) **	N019847	002	Dec 26, 1990
		1200MG/120ML (10MG/ML) **	N019847	003	Dec 26, 1990

CIPRO IN DEXTROSE 5% IN PLASTIC CONTAINER

+	BAYER HLTHCARE	200MG/100ML **	N019857	001	Dec 26, 1990
+		400MG/200ML **	N019857	002	Dec 26, 1990

CIPRO IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

	BAYER PHARMS	200MG/100ML	N019858	001	Dec 26, 1990
--	--------------	-------------	---------	-----	--------------

CIPROFLOXACIN

	BEDFORD LABS	200MG/20ML (10MG/ML)	A076992	001	Aug 28, 2006
		400MG/40ML (10MG/ML)	A076992	002	Aug 28, 2006
		1200MG/120ML (10MG/ML)	A076993	001	Aug 28, 2006
	DR REDDYS	200MG/20ML (10MG/ML)	A077782	001	Aug 28, 2006
		400MG/40ML (10MG/ML)	A077782	002	Aug 28, 2006
	FRESENIUS KABI USA	200MG/20ML (10MG/ML)	A076484	001	Aug 28, 2006
		400MG/40ML (10MG/ML)	A076484	002	Aug 28, 2006
	HOSPIRA	200MG/20ML (10MG/ML)	A077245	001	Aug 28, 2006
		400MG/40ML (10MG/ML)	A077245	002	Aug 28, 2006

CIPROFLOXACIN IN DEXTROSE 5%

	HIKMA FARMACEUTICA	200MG/100ML	A076757	001	Apr 21, 2008
--	--------------------	-------------	---------	-----	--------------

CIPROFLOXACIN IN DEXTROSE 5% IN PLASTIC CONTAINER

	BAXTER HLTHCARE	200MG/100ML	A077888	001	Mar 18, 2008
		400MG/200ML	A077888	002	Mar 18, 2008
	BAXTER HLTHCARE CORP	200MG/100ML	A078024	001	Mar 18, 2008
		400MG/200ML	A078024	002	Mar 18, 2008
	BEDFORD	200MG/100ML	A078114	001	Mar 18, 2008
		400MG/200ML	A078114	002	Mar 18, 2008
	TEVA PHARMS	200MG/100ML	A077138	001	Mar 18, 2008
		400MG/200ML	A077138	002	Mar 18, 2008

INJECTABLE, SUSPENSION; OTIC

OTIPRIO

+	ALK ABELLO	6% (60MG/ML)	N207986	001	Dec 10, 2015
---	------------	--------------	---------	-----	--------------

CIPROFLOXACIN HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

CIPROFLOXACIN HYDROCHLORIDE

	AMRING PHARMS	EQ 0.3% BASE	A078598	001	Jan 16, 2008
	RENOVA PHARMS	EQ 0.3% BASE	A076555	001	Dec 11, 2008
	THE J MOLNER	EQ 0.3% BASE	A076754	001	Jun 09, 2004

TABLET; ORAL

CIPRO

+	BAYER HLTHCARE	EQ 750MG BASE **	N019537	004	Oct 22, 1987
---	----------------	------------------	---------	-----	--------------

CIPROFLOXACIN HYDROCHLORIDE

	BARR	EQ 250MG BASE	A074124	001	Jun 09, 2004
		EQ 500MG BASE	A074124	002	Jun 09, 2004
		EQ 750MG BASE	A074124	003	Jun 09, 2004
	NATCO	EQ 250MG BASE	A075685	002	Jun 09, 2004
		EQ 500MG BASE	A075685	003	Jun 09, 2004
		EQ 750MG BASE	A075685	001	Jun 09, 2004
	NOSTRUM LABS	EQ 250MG BASE	A076138	001	Jun 09, 2004
		EQ 500MG BASE	A076138	002	Jun 09, 2004
		EQ 750MG BASE	A076138	003	Jun 09, 2004
	PLIVA	EQ 250MG BASE	A076426	002	Jun 15, 2005
		EQ 500MG BASE	A076426	003	Jun 15, 2005
		EQ 750MG BASE	A076426	004	Jun 15, 2005
	RISING	EQ 250MG BASE	A075817	002	Jun 09, 2004
		EQ 750MG BASE	A075817	004	Jun 09, 2004
	SUN PHARM INDS LTD	EQ 250MG BASE	A075747	001	Jun 09, 2004
		EQ 500MG BASE	A075747	002	Jun 09, 2004
		EQ 750MG BASE	A075747	003	Jun 09, 2004
	TARO	EQ 250MG BASE	A076912	002	Oct 06, 2004
		EQ 500MG BASE	A076912	003	Oct 06, 2004
		EQ 750MG BASE	A076912	004	Oct 06, 2004
	TEVA	EQ 250MG BASE	A076136	001	Jun 09, 2004
		EQ 500MG BASE	A076136	002	Jun 09, 2004
		EQ 750MG BASE	A076136	003	Jun 09, 2004

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CIPROFLOXACIN HYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL

PROQUIN XR

DEPOMED INC

EQ 500MG BASE

N021744 001 May 19, 2005

CIPROFLOXACIN; CIPROFLOXACIN HYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL

CIPRO XR

BAYER HLTHCARE

212.6MG;EQ 287.5MG BASE **

N021473 001 Dec 13, 2002

425.2MG;EQ 574.9MG BASE **

N021473 002 Aug 28, 2003

CIPROFLOXACIN EXTENDED RELEASE

ANI PHARMS

212.6MG;EQ 287.5MG BASE

A077809 002 Nov 30, 2010

425.2MG;EQ 574.9MG BASE

A077809 001 Nov 30, 2010

DR REDDYS LABS LTD

212.6MG;EQ 287.5MG BASE

A077701 002 Oct 31, 2007

ENDO OPERATIONS

212.6MG;EQ 287.5MG BASE

A078166 002 Nov 27, 2007

425.2MG;EQ 574.9MG BASE

A078166 001 Nov 27, 2007

FOSUN PHARMA

212.6MG;EQ 287.5MG BASE

A078712 001 Dec 11, 2007

RISING

212.6MG;EQ 287.5MG BASE

A078183 001 Mar 22, 2007

425.2MG;EQ 574.9MG BASE

A078183 002 Mar 22, 2007

CISAPRIDE MONOHYDRATE

SUSPENSION;ORAL

PROPULSID

JANSSEN PHARMS

EQ 1MG BASE/ML

N020398 001 Sep 15, 1995

TABLET;ORAL

PROPULSID

JANSSEN PHARMS

EQ 10MG BASE

N020210 001 Jul 29, 1993

EQ 20MG BASE

N020210 002 Dec 23, 1993

TABLET, ORALLY DISINTEGRATING;ORAL

PROPULSID QUICKSOLV

JANSSEN PHARMA

EQ 20MG BASE

N020767 001 Nov 07, 1997

CISATRACURIUM BESYLATE

INJECTABLE;INJECTION

CISATRACURIUM BESYLATE

ACCORD HLTHCARE

EQ 2MG BASE/ML

A205873 001 Jun 16, 2017

PIRAMAL

EQ 2MG BASE/ML

A212432 001 Mar 11, 2022

EQ 10MG BASE/ML

A212432 002 Mar 11, 2022

ZYDUS PHARMS

EQ 2MG BASE/ML

A213527 001 Aug 31, 2020

CISATRACURIUM BESYLATE PRESERVATIVE FREE

ACCORD HLTHCARE

EQ 2MG BASE/ML

A205872 001 Jun 16, 2017

EQ 10MG BASE/ML

A205872 002 Jun 16, 2017

NIMBEX

+ ABBVIE

EQ 2MG BASE/ML **

N020551 001 Dec 15, 1995

NIMBEX PRESERVATIVE FREE

+ ABBVIE

EQ 2MG BASE/ML **

N020551 003 Dec 15, 1995

+

EQ 10MG BASE/ML **

N020551 002 Dec 15, 1995

CISPLATIN

INJECTABLE;INJECTION

CISPLATIN

BEDFORD

10MG/VIAL

A074713 001 Nov 14, 2000

50MG/VIAL

A074713 002 Nov 14, 2000

NORVIUM BIOSCIENCE

1MG/ML

A091062 001 Apr 18, 2012

TEVA PHARMS USA

1MG/ML

A074814 001 May 16, 2000

PLATINOL

+ HQ SPCLT PHARMA

10MG/VIAL

N018057 001

PLATINOL-AQ

+ HQ SPCLT PHARMA

0.5MG/ML

N018057 003 Jul 18, 1984

CITALOPRAM HYDROBROMIDE

CAPSULE;ORAL

CITALOPRAM HYDROBROMIDE

NORVIUM BIOSCIENCE

EQ 10MG BASE

A077668 001 Feb 28, 2007

EQ 20MG BASE

A077668 002 Feb 28, 2007

EQ 40MG BASE

A077668 003 Feb 28, 2007

SOLUTION;ORAL

CELEXA

+ FOREST LABS

EQ 10MG BASE/5ML **

N021046 001 Dec 22, 1999

CITALOPRAM HYDROBROMIDE

PHARM ASSOC

EQ 10MG BASE/5ML

A077601 001 Nov 15, 2005

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CITALOPRAM HYDROBROMIDE

TABLET; ORAL

CELEXA

ABBVIE	EQ 60MG BASE	N020822 004	Jul 17, 1998
CITALOPRAM HYDROBROMIDE			
CHARTWELL RX	EQ 10MG BASE	A077040 001	Aug 17, 2005
	EQ 20MG BASE	A077040 002	Aug 17, 2005
	EQ 40MG BASE	A077040 003	Aug 17, 2005
COSETTE	EQ 10MG BASE	A077048 001	Nov 16, 2004
	EQ 20MG BASE	A077048 002	Nov 16, 2004
	EQ 40MG BASE	A077048 003	Nov 16, 2004
EPIC PHARMA LLC	EQ 10MG BASE	A077036 001	Oct 28, 2004
	EQ 20MG BASE	A077036 002	Oct 28, 2004
	EQ 40MG BASE	A077036 003	Oct 28, 2004
FOSUN PHARMA	EQ 10MG BASE	A077035 001	Oct 28, 2004
	EQ 20MG BASE	A077035 002	Oct 28, 2004
	EQ 40MG BASE	A077035 003	Oct 28, 2004
HERITAGE PHARMA	EQ 10MG BASE	A077033 001	Oct 28, 2004
	EQ 10MG BASE	A077034 001	Jun 30, 2005
	EQ 10MG BASE	A077213 001	Mar 31, 2006
	EQ 10MG BASE	A077232 001	Oct 31, 2005
	EQ 20MG BASE	A077033 002	Oct 28, 2004
	EQ 20MG BASE	A077034 002	Jun 30, 2005
	EQ 20MG BASE	A077213 002	Mar 31, 2006
	EQ 20MG BASE	A077232 002	Oct 31, 2005
	EQ 40MG BASE	A077033 003	Oct 28, 2004
	EQ 40MG BASE	A077034 003	Jun 30, 2005
	EQ 40MG BASE	A077213 003	Mar 31, 2006
	EQ 40MG BASE	A077232 003	Oct 31, 2005
JUBILANT GENERICS	EQ 10MG BASE	A205407 001	Dec 23, 2015
	EQ 20MG BASE	A205407 002	Dec 23, 2015
	EQ 40MG BASE	A205407 003	Dec 23, 2015
NATCO PHARMA LTD	EQ 20MG BASE	A077141 002	Apr 10, 2008
	EQ 40MG BASE	A077141 001	Apr 10, 2008
NORVIUM BIOSCIENCE	EQ 10MG BASE	A077037 001	Nov 05, 2004
	EQ 10MG BASE	A077039 001	Feb 03, 2005
	EQ 20MG BASE	A077037 002	Nov 05, 2004
	EQ 20MG BASE	A077039 002	Feb 03, 2005
	EQ 40MG BASE	A077037 003	Nov 05, 2004
	EQ 40MG BASE	A077039 003	Feb 03, 2005
ROXANE	EQ 10MG BASE	A077041 001	Nov 23, 2004
	EQ 20MG BASE	A077041 002	Nov 23, 2004
	EQ 40MG BASE	A077041 003	Nov 23, 2004
SUN PHARM INDS INC	EQ 10MG BASE	A077032 001	Nov 12, 2004
	EQ 20MG BASE	A077032 002	Nov 12, 2004
	EQ 40MG BASE	A077032 003	Nov 12, 2004
SUN PHARM INDUSTRIES	EQ 10MG BASE	A077052 001	Jul 03, 2006
	EQ 20MG BASE	A077052 002	Jul 03, 2006
	EQ 40MG BASE	A077052 003	Jul 03, 2006
TARO	EQ 10MG BASE	A077278 001	Mar 22, 2006
	EQ 20MG BASE	A077278 002	Mar 22, 2006
	EQ 40MG BASE	A077278 003	Mar 22, 2006

TABLET, ORALLY DISINTEGRATING; ORAL

CITALOPRAM HYDROBROMIDE

+	BIOVAIL LABS INTL	EQ 10MG BASE	N021763 001	Dec 20, 2005
+		EQ 20MG BASE	N021763 002	Dec 20, 2005
+		EQ 40MG BASE	N021763 003	Dec 20, 2005

CITRIC ACID; MAGNESIUM OXIDE; SODIUM CARBONATE

SOLUTION; IRRIGATION

IRRIGATING SOLUTION G IN PLASTIC CONTAINER

BAXTER HLTHCARE	3.24GM/100ML; 380MG/100ML; 430MG/100ML	N018519 001	Jun 22, 1982
UROLOGIC G IN PLASTIC CONTAINER			
HOSPIRA	3.24GM/100ML; 380MG/100ML; 430MG/100ML	N018904 001	May 27, 1983

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CITRIC ACID; MAGNESIUM OXIDE; SODIUM PICOSULFATE

FOR SOLUTION;ORAL

PREPOPIK

+ FERRING PHARMS INC 12GM/PACKET;3.5GM/PACKET;10MG/PACKET ** N202535 001 Jul 16, 2012

CLADRIBINE

INJECTABLE;INJECTION

CLADRIBINE

NORVIUM BIOSCIENCE 1MG/ML A200510 001 Oct 06, 2011

LEUSTATIN

+ JANSSEN PHARMS 1MG/ML ** N020229 001 Feb 26, 1993

CLARITHROMYCIN

FOR SUSPENSION;ORAL

BIAXIN

+ ABBVIE 125MG/5ML N050698 001 Dec 23, 1993

187MG/5ML N050698 003 Sep 30, 1998

+ 250MG/5ML N050698 002 Dec 23, 1993

CLARITHROMYCIN

SUN PHARM INDS LTD 125MG/5ML A065382 001 Aug 30, 2007

250MG/5ML A065382 002 Aug 30, 2007

TABLET;ORAL

BIAXIN

+ ABBVIE 250MG ** N050662 001 Oct 31, 1991

+ 500MG ** N050662 002 Oct 31, 1991

CLARITHROMYCIN

AJANTA PHARMA LTD 250MG A206714 001 Apr 25, 2019

500MG A206714 002 Apr 25, 2019

HIKMA 250MG A065178 002 May 25, 2004

500MG A065178 001 May 25, 2004

IVAX SUB TEVA PHARMS 250MG A065137 001 May 31, 2005

500MG A065137 002 May 31, 2005

NORVIUM BIOSCIENCE 250MG A065195 001 Mar 11, 2005

500MG A065195 002 Mar 11, 2005

SUN PHARM INDS LTD 250MG A065174 001 Sep 24, 2004

500MG A065174 002 Sep 24, 2004

TEVA 250MG A065155 001 May 31, 2005

500MG A065155 002 May 31, 2005

WOCKHARDT 250MG A065266 001 May 31, 2006

500MG A065266 002 May 31, 2006

TABLET, EXTENDED RELEASE;ORAL

BIAXIN XL

+ ABBVIE 500MG ** N050775 001 Mar 03, 2000

CLARITHROMYCIN

ANI PHARMS 500MG A065250 001 Aug 25, 2005

LUPIN LTD 500MG A202532 001 Sep 15, 2015

RANBAXY 1GM A065210 001 Jan 26, 2005

CLAVULANATE POTASSIUM; TICARCILLIN DISODIUM

INJECTABLE;INJECTION

TIMENTIN

GLAXOSMITHKLINE EQ 100MG BASE/VIAL;EQ 3GM BASE/VIAL A062691 001 Dec 19, 1986

EQ 100MG BASE/VIAL;EQ 3GM BASE/VIAL N050590 001 Apr 01, 1985

EQ 200MG BASE/VIAL;EQ 3GM BASE/VIAL N050590 002 Apr 01, 1985

EQ 1GM BASE/VIAL;EQ 30GM BASE/VIAL N050590 003 Aug 18, 1987

TIMENTIN IN PLASTIC CONTAINER

GLAXOSMITHKLINE EQ 100MG BASE/100ML;EQ 3GM BASE/100ML N050658 001 Dec 15, 1989

CLEMASTINE FUMARATE

SYRUP;ORAL

CLEMASTINE FUMARATE

ACTAVIS MID ATLANTIC EQ 0.5MG BASE/5ML A074075 001 Oct 31, 1993

APOTEX INC EQ 0.5MG BASE/5ML A075703 001 Nov 27, 2000

TEVA PHARMS EQ 0.5MG BASE/5ML A073095 001 Apr 21, 1992

WOCKHARDT BIO AG EQ 0.5MG BASE/5ML A074863 001 Mar 13, 1998

TAVIST

+ NOVARTIS EQ 0.5MG BASE/5ML ** N018675 001 Jun 28, 1985

TABLET;ORAL

CLEMASTINE FUMARATE

ANI PHARMS 1.34MG A073282 001 Jan 31, 1992

1.34MG A073282 002 Dec 03, 1992

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CLEMASTINE FUMARATE

TABLET; ORAL

CLEMASTINE FUMARATE

L PERRIGO CO	1.34MG	A074512	001	Nov 22, 1995
OMNIVIUM PHARMS	2.68MG	A073459	001	Oct 31, 1993
PLD ACQUISITIONS LLC	1.34MG	A073458	001	Oct 31, 1993

TAVIST

+ NOVARTIS	2.68MG **	N017661	001	
------------	-----------	---------	-----	--

TAVIST-1

+ HALEON US HOLDINGS	1.34MG **	N020925	001	Aug 21, 1992
NOVARTIS	1.34MG	N017661	002	
	1.34MG	N017661	003	Aug 21, 1992

CLEVIDIPINE

EMULSION; INTRAVENOUS

CLEVIPREX

+ CHIESI	125MG/250ML (0.5MG/ML)	N022156	003	Nov 08, 2013
----------	------------------------	---------	-----	--------------

CLIDINIUM BROMIDE

CAPSULE; ORAL

QUARZAN

ROCHE	2.5MG	N010355	001	
	5MG	N010355	002	

CLINDAMYCIN HYDROCHLORIDE

CAPSULE; ORAL

CLEOCIN

PHARMACIA AND UPJOHN	EQ 75MG BASE	A061809	001	
	EQ 150MG BASE	A061809	002	

CLINDAMYCIN HYDROCHLORIDE

CHARTWELL MOLECULAR	EQ 150MG BASE	A065243	003	Aug 12, 2005
	EQ 300MG BASE	A065243	001	Aug 12, 2005
COSETTE	EQ 150MG BASE	A063029	001	Sep 20, 1989
	EQ 300MG BASE	A063029	002	Aug 05, 2005
NORVIUM BIOSCIENCE	EQ 75MG BASE	A091225	001	May 31, 2011
	EQ 150MG BASE	A091225	002	May 31, 2011
	EQ 300MG BASE	A091225	003	May 31, 2011
TEVA	EQ 75MG BASE	A063027	001	Sep 20, 1989
WATSON LABS	EQ 75MG BASE	A063082	001	Jul 31, 1991
	EQ 150MG BASE	A063083	001	Jul 31, 1991
	EQ 300MG BASE	A063083	002	Mar 18, 2003

CLINDAMYCIN PALMITATE HYDROCHLORIDE

FOR SOLUTION; ORAL

CLEOCIN

PHARMACIA AND UPJOHN	EQ 75MG BASE/5ML **	A061827	001	
----------------------	---------------------	---------	-----	--

CLINDAMYCIN PALMITATE HYDROCHLORIDE

EXTROVIS	EQ 75MG BASE/5ML	A203063	001	May 25, 2016
----------	------------------	---------	-----	--------------

CLINDAMYCIN PHOSPHATE

AEROSOL, FOAM; TOPICAL

EVOCLIN

+ NORVIUM BIOSCIENCE	1% **	N050801	001	Oct 22, 2004
----------------------	-------	---------	-----	--------------

CREAM; VAGINAL

CLEOCIN

PFIZER	EQ 2% BASE	N050680	001	Aug 11, 1992
--------	------------	---------	-----	--------------

GEL; TOPICAL

CLINDAMYCIN PHOSPHATE

AMNEAL	EQ 1% BASE	A215219	001	Nov 03, 2022
--------	------------	---------	-----	--------------

INJECTABLE; INJECTION

CLEOCIN PHOSPHATE

PHARMACIA AND UPJOHN	EQ 150MG BASE/ML	A061839	001	
----------------------	------------------	---------	-----	--

CLEOCIN PHOSPHATE IN DEXTROSE 5% IN PLASTIC CONTAINER

+ PFIZER	EQ 6MG BASE/ML **	N050639	001	Aug 30, 1989
	EQ 12MG BASE/ML **	N050639	002	Aug 30, 1989
	EQ 18MG BASE/ML **	N050639	003	Apr 10, 1991

CLINDAMYCIN PHOSPHATE

ABRAXIS PHARM	EQ 150MG BASE/ML	A062747	001	Jun 03, 1988
ALMAJECT	EQ 150MG BASE/ML	A062801	001	Jul 24, 1987
BEDFORD	EQ 150MG BASE/ML	A063163	001	Jun 30, 1994
BRISTOL MYERS SQUIBB	EQ 150MG BASE/ML	A062908	001	Feb 01, 1989
HIKMA	EQ 150MG BASE/ML	A062806	001	Oct 15, 1987

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CLINDAMYCIN PHOSPHATE

INJECTABLE; INJECTION

CLINDAMYCIN PHOSPHATE

	EQ 150MG BASE/ML	A062953	001	Apr 21, 1988
	EQ 150MG BASE/ML	A063068	001	Aug 28, 1989
IGI LABS INC	EQ 150MG BASE/ML	A062928	001	Feb 13, 1989
LOCH	EQ 150MG BASE/ML	A062905	001	May 09, 1988
MARSAM PHARMS LLC	EQ 150MG BASE/ML	A062913	001	Oct 20, 1988
RISING	EQ 150MG BASE/ML	A204748	001	Oct 10, 2017
	EQ 150MG BASE/ML	A204749	001	Oct 10, 2017
SOLOPAK	EQ 150MG BASE/ML	A062819	001	Mar 15, 1988
	EQ 150MG BASE/ML	A062852	001	Mar 17, 1988
TEVA PARENTERAL	EQ 150MG BASE/ML	A063041	001	Dec 29, 1989
	EQ 150MG BASE/ML	A063282	001	May 29, 1992
WATSON LABS	EQ 150MG BASE/ML	A062900	001	Jun 08, 1988
	EQ 150MG BASE/ML	A063079	001	Mar 05, 1990
CLINDAMYCIN PHOSPHATE IN DEXTROSE 5%				
ABRAXIS PHARM	EQ 12MG BASE/ML	N050636	001	Dec 22, 1989
CLINDAMYCIN PHOSPHATE IN DEXTROSE 5% IN PLASTIC CONTAINER				
ABBOTT LABS	EQ 6MG BASE/ML	A065027	001	Jun 29, 2001
	EQ 12MG BASE/ML	A065027	002	Jun 29, 2001
	EQ 18MG BASE/ML	A065027	003	Jun 29, 2001
BAXTER HLTHCARE	EQ 6MG BASE/ML	N050648	001	Dec 29, 1989
	EQ 12MG BASE/ML	N050648	002	Dec 29, 1989
	EQ 900MG BASE/100ML	N050648	003	Dec 29, 1989

SOLUTION; TOPICAL

CLEOCIN T

+	PFIZER	EQ 1% BASE **	N050537	001	
	PHARMACIA AND UPJOHN	EQ 1% BASE	A062363	001	Feb 08, 1982

CLINDAMYCIN PHOSPHATE

BOCA PHARMA LLC	EQ 1% BASE	A062944	001	Jan 11, 1989
ENDO OPERATIONS	EQ 1% BASE	A203343	001	May 29, 2015
FOUGERA PHARMS	EQ 1% BASE	A065254	001	Feb 14, 2006
G AND W LABS INC	EQ 1% BASE	A062811	001	Sep 01, 1988
NOVAST LABS	EQ 1% BASE	A064108	001	Sep 27, 1996
PAI HOLDINGS PHARM	EQ 1% BASE	A206945	001	Dec 30, 2016
VINTAGE PHARMS	EQ 1% BASE	A062930	001	Jun 28, 1989
XTRTRIUM LABS INC	EQ 1% BASE	A063304	001	Jul 15, 1997

SWAB; TOPICAL

CLEOCIN

+	PFIZER	EQ 1% BASE	N050537	002	Feb 22, 1994
---	--------	------------	---------	-----	--------------

CLIOQUINOL; NYSTATIN

OINTMENT; TOPICAL

NYSTAFORM

BAYER PHARMS	10MG/GM; 100,000 UNITS/GM	N050235	001	
--------------	---------------------------	---------	-----	--

CLOBAZAM

SUSPENSION; ORAL

CLOBAZAM

ACCORD HLTHCARE	2.5MG/ML	A216008	001	Sep 06, 2022
HIKMA	2.5MG/ML	A209715	001	Oct 22, 2018
NORVIUM BIOSCIENCE	2.5MG/ML	A211259	001	Oct 22, 2018
TEVA PHARMS USA	2.5MG/ML	A211032	001	Jan 31, 2020
UPSHER SMITH LABS	2.5MG/ML	A210569	001	Oct 22, 2018
VISTAPHARM	2.5MG/ML	A210746	001	Jul 10, 2019

TABLET; ORAL

CLOBAZAM

ACCORD HLTHCARE	10MG	A212398	001	May 23, 2019
	20MG	A212398	002	May 23, 2019
ALLIED	5MG	A209308	003	Oct 19, 2021
	20MG	A209308	002	Oct 22, 2018
ANDA REPOSITORY	10MG	A211959	001	Dec 09, 2020
	20MG	A211959	002	Dec 09, 2020
APOTEX	10MG	A209853	001	Jun 09, 2020
	20MG	A209853	002	Jun 09, 2020
CHARTWELL MOLECULAR	10MG	A212092	001	Oct 30, 2019
	20MG	A212092	002	Oct 30, 2019
HIKMA	10MG	A208785	001	Oct 22, 2018
	20MG	A208785	002	Oct 22, 2018

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CLOBAZAM

TABLET; ORAL

CLOBAZAM

TARO

10MG

A209440 001 Oct 22, 2018

20MG

A209440 002 Oct 22, 2018

ONFI

+ LUNDBECK PHARMS LLC 5MG **

N202067 001 Oct 21, 2011

CLOBETASOL PROPIONATE

AEROSOL, FOAM; TOPICAL

CLOBETASOL PROPIONATE

NOVAST LABS

0.05%

A206805 001 Jul 31, 2017

OLUX

+ NORVIUM BIOSCIENCE 0.05%

N021142 001 May 26, 2000

OLUX E

+ NORVIUM BIOSCIENCE 0.05% **

N022013 001 Jan 12, 2007

CREAM; TOPICAL

CLOBETASOL PROPIONATE

CHARTWELL RX

0.05%

A211207 001 Mar 26, 2021

COSETTE

0.05%

A074139 001 Aug 03, 1994

PAI HOLDINGS PHARM

0.05%

A209974 001 Apr 17, 2018

TEVA PHARMS USA

0.05%

A074087 001 Feb 16, 1994

CLOBETASOL PROPIONATE (EMOLLIENT)

ANI PHARMS

0.05%

A075733 001 Aug 22, 2001

CORMAX

HIKMA

0.05%

A074220 001 May 16, 1997

EMBELINE E

HIKMA

0.05%

A075325 001 Dec 24, 1998

TEMOVATE

+ FOUGERA PHARMS 0.05% **

N019322 001 Dec 27, 1985

TEMOVATE E

+ FOUGERA PHARMS 0.05% **

N020340 001 Jun 17, 1994

GEL; TOPICAL

CLOBETASOL PROPIONATE

PAI HOLDINGS PHARM

0.05%

A208881 001 Mar 06, 2017

EMBELINE

HIKMA

0.05%

A076141 001 Apr 12, 2002

TEMOVATE

+ FOUGERA PHARMS 0.05% **

N020337 001 Apr 29, 1994

LOTION; TOPICAL

CLOBETASOL PROPIONATE

EPIC PHARMA LLC

0.05%

A211348 001 Oct 26, 2018

PAI HOLDINGS PHARM

0.05%

A208667 001 Nov 29, 2016

IMPEKLO

+ MYLAN 0.05%

N213691 001 May 19, 2020

OINTMENT; TOPICAL

CLOBETASOL PROPIONATE

ACTAVIS MID ATLANTIC

0.05%

A074128 001 Aug 03, 1994

AMNEAL

0.05%

A210551 001 Aug 21, 2018

PAI HOLDINGS PHARM

0.05%

A208589 001 Jan 23, 2019

TORRENT

0.05%

A212926 001 Oct 25, 2019

EMBELINE

HIKMA

0.05%

A074221 001 Mar 31, 1995

TEMOVATE

+ FOUGERA PHARMS 0.05% **

N019323 001 Dec 27, 1985

SHAMPOO; TOPICAL

CLOBETASOL PROPIONATE

ACTAVIS MID ATLANTIC

0.05%

A078854 001 Jun 07, 2011

HIKMA

0.05%

A209871 001 Oct 27, 2017

SOLUTION; TOPICAL

CLOBETASOL PROPIONATE

COSETTE

0.05%

A074331 001 Dec 15, 1995

PRINSTON INC

0.05%

A213139 001 Feb 08, 2021

WOCKHARDT BIO AG

0.05%

A075205 001 Nov 13, 1998

EMBELINE

HIKMA

0.05%

A074222 001 Dec 06, 1995

TEMOVATE

+ FOUGERA PHARMS 0.05% **

N019966 001 Feb 22, 1990

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CLOBETASOL PROPIONATE

SPRAY;TOPICAL

CLOBETASOL PROPIONATE

APOTEX	0.05%	A210446	001	Apr 17, 2018
SCIEGEN PHARMS INC	0.05%	A207218	001	Apr 28, 2017

CLOFARABINE

SOLUTION;INTRAVENOUS

CLOFARABINE

ABON PHARMS LLC	20MG/20ML (1MG/ML)	A204029	001	May 09, 2017
EUGIA PHARMA	20MG/20ML (1MG/ML)	A212457	001	Oct 03, 2022
HOSPIRA	20MG/20ML (1MG/ML)	A210283	001	Dec 27, 2018
NOVAST LABS	20MG/20ML (1MG/ML)	A210270	001	Sep 14, 2018

CLOFAZIMINE

CAPSULE;ORAL

LAMPRENE

+ NOVARTIS	50MG	N019500	002	Dec 15, 1986
	100MG	N019500	001	Dec 15, 1986

CLOFIBRATE

CAPSULE;ORAL

ATROMID-S

WYETH AYERST	500MG	N016099	002	
--------------	-------	---------	-----	--

CLOFIBRATE

BANNER PHARMACAPS	500MG	A073396	001	Mar 20, 1992
SANDOZ	500MG	A072191	001	May 02, 1988
TEVA	500MG	A072600	001	Jul 25, 1991
USL PHARMA	500MG	A070531	001	Jun 16, 1986
WATSON LABS	500MG	A071603	001	Sep 18, 1987

CLOMIPHENE CITRATE

TABLET;ORAL

CLOMID

+ SANOFI AVENTIS US	50MG **	N016131	002	
---------------------	---------	---------	-----	--

MILOPHENE

GRANATA BIO	50MG	A072196	001	Dec 20, 1988
-------------	------	---------	-----	--------------

SEROPHENE

EMD SERONO	50MG	N018361	001	Mar 22, 1982
------------	------	---------	-----	--------------

CLOMIPRAMINE HYDROCHLORIDE

CAPSULE;ORAL

CLOMIPRAMINE HYDROCHLORIDE

AJANTA PHARMA LTD	25MG	A213897	001	Oct 02, 2020
	50MG	A213897	002	Oct 02, 2020
	75MG	A213897	003	Oct 02, 2020
AMNEAL PHARMS CO	25MG	A208632	001	Oct 31, 2018
	50MG	A208632	002	Oct 31, 2018
	75MG	A208632	003	Oct 31, 2018
MYLAN	25MG	A074947	001	Apr 30, 1998
	50MG	A074947	002	Apr 30, 1998
	75MG	A074947	003	Apr 30, 1998
RK PHARMA	25MG	A213221	001	Jun 22, 2020
	50MG	A213221	002	Jun 22, 2020
	75MG	A213221	003	Jun 22, 2020
TEVA	25MG	A074849	001	Apr 04, 1997
	25MG	A074958	001	Aug 26, 1997
	50MG	A074849	002	Apr 04, 1997
	50MG	A074958	002	Aug 26, 1997
	75MG	A074849	003	Apr 04, 1997
	75MG	A074958	003	Aug 26, 1997
WATSON LABS	25MG	A074600	001	Nov 27, 1996
	25MG	A074751	001	Sep 30, 1998
	50MG	A074600	002	Nov 27, 1996
	50MG	A074751	002	Sep 30, 1998
	75MG	A074600	003	Nov 27, 1996
	75MG	A074751	003	Sep 30, 1998

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CLONAZEPAM

TABLET; ORAL

CLONAZEPAM

CHARTWELL RX	0.5MG	A074925 001	Sep 30, 1997
	1MG	A074925 002	Sep 30, 1997
	2MG	A074925 003	Sep 30, 1997
NORVIUM BIOSCIENCE	0.5MG	A074940 001	Oct 30, 1997
	1MG	A074940 002	Oct 30, 1997
	2MG	A074940 003	Oct 30, 1997
SUN PHARM INDS INC	0.5MG	A075423 001	Apr 27, 2001
	1MG	A075423 002	Apr 27, 2001
	2MG	A075423 003	Apr 27, 2001
TEVA	0.5MG	A074920 001	Aug 04, 1998
	1MG	A074920 002	Aug 04, 1998
	2MG	A074920 003	Aug 04, 1998
WATSON LABS	0.5MG	A074964 001	Dec 30, 1997
	1MG	A074964 002	Dec 30, 1997
	2MG	A074964 003	Dec 30, 1997

KLONOPIN

CHEPLAPHARM	0.125MG	N017533 005	Apr 09, 1997
	0.25MG	N017533 006	Apr 09, 1997

TABLET, ORALLY DISINTEGRATING; ORAL

KLONOPIN RAPIDLY DISINTEGRATING

+ ROCHE	0.125MG **	N020813 001	Dec 23, 1997
+	0.25MG **	N020813 002	Dec 23, 1997
+	0.5MG **	N020813 003	Dec 23, 1997
+	1MG **	N020813 004	Dec 23, 1997
+	2MG **	N020813 005	Dec 23, 1997

CLONIDINE

SUSPENSION, EXTENDED RELEASE; ORAL

CLONIDINE

TRIS PHARMA INC	EQ 0.09MG BASE/ML	N022499 001	Dec 03, 2009
-----------------	-------------------	-------------	--------------

TABLET, EXTENDED RELEASE; ORAL

NEXICLON XR

ATHENA	EQ 0.26MG BASE	N022500 002	Dec 03, 2009
--------	----------------	-------------	--------------

CLONIDINE HYDROCHLORIDE

INJECTABLE; INJECTION

CLONIDINE HYDROCHLORIDE

AM REGENT	1MG/10ML (0.1MG/ML)	A091104 001	Oct 08, 2009
	5MG/10ML (0.5MG/ML)	A091104 002	Oct 08, 2009

DURACLON

+ MYLAN INSTITUTIONAL	5MG/10ML (0.5MG/ML) **	N020615 002	Apr 27, 1999
-----------------------	------------------------	-------------	--------------

TABLET; ORAL

CATAPRES

+ BOEHRINGER INGELHEIM	0.1MG **	N017407 001	
+	0.2MG **	N017407 002	
+	0.3MG **	N017407 003	

CLONIDINE HYDROCHLORIDE

AM THERAP	0.1MG	A070881 001	Jul 08, 1986
	0.2MG	A070882 001	Jul 08, 1986
	0.3MG	A070883 001	Jul 08, 1986
DURAMED PHARMS BARR	0.1MG	A071103 001	Aug 14, 1986
	0.2MG	A071102 001	Aug 14, 1986
	0.3MG	A071101 001	Aug 14, 1986
INTERPHARM	0.1MG	A071252 001	Oct 01, 1986
	0.2MG	A071253 001	Oct 01, 1986
	0.3MG	A071254 001	Oct 01, 1986
PAR PHARM	0.1MG	A070461 001	Jul 08, 1986
	0.2MG	A070460 001	Jul 08, 1986
	0.3MG	A070459 001	Jul 08, 1986
RISING	0.1MG	A070317 002	Jul 09, 1987
	0.2MG	A070317 003	Jun 09, 1987
	0.3MG	A070317 001	Jun 09, 1987
SUN PHARM INDS INC	0.1MG	A090329 001	Jul 03, 2014
	0.2MG	A090329 002	Jul 03, 2014
	0.3MG	A090329 003	Jul 03, 2014
TEVA	0.1MG	A070747 001	Jul 08, 1986
	0.2MG	A070702 001	Jul 08, 1986

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CLONIDINE HYDROCHLORIDE

TABLET;ORAL

CLONIDINE HYDROCHLORIDE

	0.3MG	A070659 001	Jul 08, 1986
WARNER CHILCOTT	0.1MG	A072138 001	Jun 13, 1988
	0.2MG	A072139 001	Jun 13, 1988
	0.3MG	A072140 001	Jun 13, 1988
WATSON LABS	0.1MG	A070395 001	Mar 23, 1987
	0.1MG	A070965 001	Jul 08, 1986
	0.2MG	A070396 001	Mar 23, 1987
	0.2MG	A070964 001	Jul 08, 1986
	0.3MG	A070397 001	Mar 23, 1987
	0.3MG	A070963 001	Jul 08, 1986

TABLET, EXTENDED RELEASE;ORAL

CLONIDINE HYDROCHLORIDE

ACTAVIS ELIZABETH	0.1MG	A202792 001	May 15, 2015
	0.2MG	A202792 002	May 15, 2015
	0.2MG	A203320 002	May 15, 2015
AMNEAL PHARMS NY	0.1MG	A210052 001	Nov 20, 2017
DR REDDYS LABS SA	0.1MG	A210680 001	Apr 30, 2018
ENDO OPERATIONS	0.1MG	A202983 001	Apr 02, 2014
	0.1MG	A202984 001	Sep 30, 2013
	0.2MG	A202983 002	Apr 02, 2014
	0.2MG	A202984 002	Sep 30, 2013
SOMERSET THERAPS LLC	0.1MG	A211433 001	Oct 12, 2018
JENLOGA			
+ CONCORDIA PHARMS INC	0.1MG **	N022331 001	Sep 30, 2009
+	0.2MG **	N022331 002	May 25, 2010
KAPVAY			
+ CONCORDIA PHARMS INC	0.1MG	N022331 003	Sep 28, 2010
+	0.2MG **	N022331 004	Sep 28, 2010

CLOPIDOGREL BISULFATE

TABLET;ORAL

CLOPIDOGREL BISULFATE

ACCORD HLTHCARE	EQ 75MG BASE	A202925 001	Mar 27, 2013
	EQ 300MG BASE	A202925 002	Mar 27, 2013
ACTAVIS TOTOWA	EQ 75MG BASE	A090307 001	May 28, 2013
ANI PHARMS	EQ 300MG BASE	A090625 001	May 17, 2012
	EQ 300MG BASE	A091216 001	May 17, 2012
CHARTWELL RX	EQ 75MG BASE	A202266 001	Aug 14, 2012
	EQ 300MG BASE	A202266 002	Nov 20, 2012
RISING	EQ 75MG BASE	A077665 001	May 17, 2012
	EQ 300MG BASE	A077665 002	May 17, 2012
SUN PHARM	EQ 75MG BASE	A090494 001	May 17, 2012
SUN PHARM INDUSTRIES	EQ 75MG BASE	A078133 001	Jun 10, 2013
ZYDUS LIFESCIENCES	EQ 75MG BASE	A201686 001	Oct 10, 2012

CLORAZEPATE DIPOTASSIUM

CAPSULE;ORAL

CLORAZEPATE DIPOTASSIUM

ABLE	3.75MG	A071777 001	Jul 14, 1987
	7.5MG	A071778 001	Jul 14, 1987
	15MG	A071779 001	Jul 14, 1987
AM THERAP	3.75MG	A071429 001	Jun 23, 1987
	7.5MG	A071430 001	Jun 23, 1987
	15MG	A071431 001	Jun 23, 1987
DASH PHARMS	3.75MG	A071509 001	Oct 19, 1987
	7.5MG	A071510 001	Oct 19, 1987
	15MG	A071511 001	Oct 19, 1987
DAVA PHARMS INC	3.75MG	A071742 001	Dec 14, 1987
	7.5MG	A071743 001	Dec 14, 1987
	15MG	A071744 001	Dec 14, 1987
GD SEARLE LLC	3.75MG	A071727 001	Dec 18, 1987
	7.5MG	A071728 001	Dec 18, 1987
	15MG	A071729 001	Dec 18, 1987
PUREPAC PHARM	3.75MG	A071924 001	Apr 25, 1988
	7.5MG	A071925 001	Apr 25, 1988
	15MG	A071926 001	Apr 25, 1988
QUANTUM PHARMICS	3.75MG	A071549 001	Sep 12, 1988

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CLORAZEPATE DIPOTASSIUM

CAPSULE;ORAL

CLORAZEPATE DIPOTASSIUM

	7.5MG	A071550 001	Sep 12, 1988
	15MG	A071522 001	Sep 12, 1988
RISING	3.75MG	A072112 002	Aug 26, 1988
	7.5MG	A072112 003	Aug 26, 1988
	15MG	A072112 001	Aug 26, 1988
USL PHARMA	3.75MG	A071242 001	Jun 23, 1987
	7.5MG	A071243 001	Jun 23, 1987
	15MG	A071244 001	Jun 23, 1987
WARNER CHILCOTT	3.75MG	A071774 001	Mar 01, 1988
	7.5MG	A071775 001	Mar 01, 1988
	15MG	A071776 001	Mar 01, 1988
WATSON LABS	3.75MG	A071878 001	Mar 15, 1988
	7.5MG	A071879 001	Mar 15, 1988
	15MG	A071860 001	Mar 15, 1988
TRANXENE			
+ AJENAT PHARMS	3.75MG **	N017105 001	
+	7.5MG **	N017105 002	
+	15MG **	N017105 003	

TABLET;ORAL

CLORAZEPATE DIPOTASSIUM

ABLE	3.75MG	A071780 001	Jun 26, 1987
	7.5MG	A071781 001	Jun 26, 1987
	15MG	A071782 001	Jun 26, 1987
AM THERAP	3.75MG	A071747 001	Jun 23, 1987
	7.5MG	A071748 001	Jun 23, 1987
	15MG	A071749 001	Jun 23, 1987
AUROLIFE PHARMA LLC	3.75MG	A072514 002	May 11, 1990
	7.5MG	A072514 003	May 11, 1990
	15MG	A072514 001	May 11, 1990
LEDERLE	3.75MG	A072013 001	Dec 15, 1987
	7.5MG	A072014 001	Dec 15, 1987
	15MG	A072015 001	Dec 15, 1987
PUREPAC PHARM	3.75MG	A072330 001	Aug 08, 1988
	7.5MG	A072331 001	Aug 08, 1988
	15MG	A072332 001	Aug 08, 1988
QUANTUM PHARMICS	3.75MG	A071730 001	Oct 26, 1987
	7.5MG	A071731 001	Oct 26, 1987
	15MG	A071702 001	Oct 26, 1987
SUN PHARM INDS LTD	3.75MG	A076911 001	Sep 29, 2004
	7.5MG	A076911 002	Sep 29, 2004
	15MG	A076911 003	Sep 29, 2004
WARNER CHILCOTT	3.75MG	A071828 001	Mar 03, 1988
	7.5MG	A071829 001	Mar 03, 1988
	15MG	A071830 001	Mar 03, 1988
WATSON LABS	3.75MG	A071852 001	Feb 09, 1988
	7.5MG	A071853 001	Feb 09, 1988
	15MG	A071854 001	Feb 09, 1988
GEN-XENE			
ALRA	3.75MG	A071787 001	Apr 26, 1988
	7.5MG	A071788 001	Apr 26, 1988
	15MG	A071789 001	Apr 26, 1988
TRANXENE			
+ AJENAT PHARMS	3.75MG **	N017105 006	
+	15MG **	N017105 008	
TRANXENE SD			
+ AJENAT PHARMS	11.25MG **	N017105 005	
+	22.5MG **	N017105 004	

CLOTRIMAZOLE

CREAM;TOPICAL

LOTRIMIN

+ SCHERING PLOUGH	1% **	N017619 001	
MYCELEX			
BAYER HEALTHCARE LLC	1%	N018183 001	

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CLOTRIMAZOLE

CREAM;VAGINAL

GYNE-LOTRIMIN

+ BAYER HEALTHCARE LLC 1% **

N018052 002 Nov 30, 1990

GYNE-LOTRIMIN 3

+ BAYER HEALTHCARE LLC 2%

N020574 001 Nov 24, 1998

MYCELEX-7

BAYER HEALTHCARE LLC 1%

N018230 002 Dec 26, 1991

CREAM, TABLET;TOPICAL, VAGINAL

GYNE-LOTRIMIN 3 COMBINATION PACK

+ BAYER HEALTHCARE LLC 1%,200MG

N020526 002 Jul 29, 1996

GYNE-LOTRIMIN COMBINATION PACK

+ BAYER HEALTHCARE LLC 1%,100MG

N020289 002 Apr 26, 1993

MYCELEX-7 COMBINATION PACK

BAYER HEALTHCARE LLC 1%,100MG

N020389 002 Jun 23, 1994

LOTION;TOPICAL

LOTRIMIN

SCHERING 1%

N018813 001 Feb 17, 1984

SOLUTION;TOPICAL

LOTRIMIN

+ SCHERING PLOUGH 1% **

N017613 001

MYCELEX

+ BAYER HLTHCARE 1% **

N018181 001

TABLET;VAGINAL

GYNE-LOTRIMIN

+ BAYER HEALTHCARE LLC 100MG

N017717 002 Nov 30, 1990

GYNE-LOTRIMIN 3

+ BAYER HEALTHCARE LLC 200MG

N020525 001 Jul 29, 1996

GYNIX

TEVA PHARMS 100MG

A073249 001 Feb 13, 1998

MYCELEX-7

BAYER HEALTHCARE LLC 100MG

N018182 002 Dec 26, 1991

MYCELEX-G

BAYER PHARMS 500MG

N019069 001 Apr 19, 1985

TROCHE/LOZENGE;ORAL

MYCELEX

+ BAYER HLTHCARE 10MG **

N018713 001 Jun 17, 1983

CLOXACILLIN SODIUM

CAPSULE;ORAL

CLOXACILLIN SODIUM

APOTHECON EQ 250MG BASE

A061452 001

EQ 500MG BASE

A061452 002

TEVA EQ 250MG BASE

A062240 001

EQ 500MG BASE

A062240 002

CLOXAPEN

GLAXOSMITHKLINE EQ 250MG BASE

A061806 001

EQ 250MG BASE

A062233 001

EQ 500MG BASE

A061806 002

EQ 500MG BASE

A062233 002

FOR SOLUTION;ORAL

CLOXACILLIN SODIUM

TEVA EQ 125MG BASE/5ML

A062268 001

EQ 125MG BASE/5ML

A062978 001 Apr 06, 1989

TEGOPEN

APOTHECON EQ 125MG BASE/5ML

A061453 001

EQ 125MG BASE/5ML

N050192 001

CLOZAPINE

TABLET;ORAL

CLOZAPINE

DR REDDYS LABS SA 25MG

A203807 001 Sep 17, 2015

50MG

A203807 003 Aug 22, 2017

100MG

A203807 002 Sep 17, 2015

200MG

A203807 004 Aug 22, 2017

MYLAN 12.5MG

A075417 003 Apr 15, 2010

PAR PHARM 25MG

A075162 001 Apr 26, 2005

100MG

A075162 002 Apr 26, 2005

SANDOZ 25MG

A074546 001 Aug 30, 1996

100MG

A074546 002 Aug 30, 1996

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CLOZAPINE

TABLET;ORAL

CLOZAPINE

ZYDUS PHARMS	25MG	A209480 001	Dec 06, 2017
	50MG	A209480 002	Dec 06, 2017
	100MG	A209480 003	Dec 06, 2017
	200MG	A209480 004	Dec 06, 2017

CLOZARIL

+ HERITAGE LIFE	50MG	N019758 003	May 20, 2019
+	200MG	N019758 004	May 20, 2019

TABLET, ORALLY DISINTEGRATING;ORAL

FAZACLO ODT

+ JAZZ	12.5MG **	N021590 004	May 30, 2007
+	25MG **	N021590 001	Feb 10, 2004
	50MG **	N021590 003	Jun 03, 2005
+	100MG **	N021590 002	Feb 10, 2004
+	150MG **	N021590 005	Jul 09, 2010
+	200MG **	N021590 006	Jul 09, 2010

COBICISTAT

TABLET;ORAL

COBICISTAT

MYLAN LABS LTD	150MG	A209986 001	Feb 07, 2024
----------------	-------	-------------	--------------

CODEINE PHOSPHATE; PHENYLEPHRINE HYDROCHLORIDE; PROMETHAZINE HYDROCHLORIDE

SYRUP;ORAL

PHENERGAN VC W/ CODEINE

+ ANI PHARMS	10MG/5ML;5MG/5ML;6.25MG/5ML **	N008306 005	Apr 02, 1984
--------------	--------------------------------	-------------	--------------

PHERAZINE VC W/ CODEINE

HALSEY	10MG/5ML;5MG/5ML;6.25MG/5ML	A088870 001	Mar 02, 1987
--------	-----------------------------	-------------	--------------

PROMETH VC W/ CODEINE

NOSTRUM LABS INC	10MG/5ML;5MG/5ML;6.25MG/5ML	A088764 001	Oct 31, 1984
------------------	-----------------------------	-------------	--------------

PROMETHAZINE HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE W/CODEINE PHOSPHATE

HIKMA	10MG/5ML;5MG/5ML;6.25MG/5ML	A040674 001	Dec 23, 2014
-------	-----------------------------	-------------	--------------

PROMETHAZINE VC W/ CODEINE

CENCI	10MG/5ML;5MG/5ML;6.25MG/5ML	A088816 001	Nov 22, 1985
-------	-----------------------------	-------------	--------------

WOCKHARDT	10MG/5ML;5MG/5ML;6.25MG/5ML	A088896 001	Jan 04, 1985
-----------	-----------------------------	-------------	--------------

CODEINE PHOSPHATE; PROMETHAZINE HYDROCHLORIDE

SYRUP;ORAL

PHENERGAN W/ CODEINE

+ ANI PHARMS	10MG/5ML;6.25MG/5ML **	N008306 004	Apr 02, 1984
--------------	------------------------	-------------	--------------

PHERAZINE W/ CODEINE

HALSEY	10MG/5ML;6.25MG/5ML	A088739 001	Dec 23, 1988
--------	---------------------	-------------	--------------

PROMETHAZINE HYDROCHLORIDE AND CODEINE PHOSPHATE

+ ACTAVIS MID ATLANTIC	10MG/5ML;6.25MG/5ML	A088763 001	Oct 31, 1984
------------------------	---------------------	-------------	--------------

HIKMA	10MG/5ML;6.25MG/5ML	A040151 001	Aug 26, 1997
-------	---------------------	-------------	--------------

PHARM ASSOC	10MG/5ML;6.25MG/5ML	A089647 001	Dec 22, 1988
-------------	---------------------	-------------	--------------

PROMETHAZINE W/ CODEINE

CENCI	10MG/5ML;6.25MG/5ML	A088814 001	Nov 22, 1985
-------	---------------------	-------------	--------------

CODEINE PHOSPHATE; PSEUDOEPHEDRINE HYDROCHLORIDE; TRIPROLIDINE HYDROCHLORIDE

SYRUP;ORAL

ACTIFED W/ CODEINE

GLAXOSMITHKLINE	10MG/5ML;30MG/5ML;1.25MG/5ML **	N012575 003	Apr 04, 1984
-----------------	---------------------------------	-------------	--------------

TRIPROLIDINE AND PSEUDOEPHEDRINE HYDROCHLORIDES W/ CODEINE

CENCI	10MG/5ML;30MG/5ML;1.25MG/5ML	A089018 001	Jul 23, 1986
-------	------------------------------	-------------	--------------

TRIPROLIDINE HYDROCHLORIDE, PSEUDOEPHEDRINE HYDROCHLORIDE AND CODEINE PHOSPHATE

WOCKHARDT	10MG/5ML;30MG/5ML;1.25MG/5ML	A088833 001	Nov 16, 1984
-----------	------------------------------	-------------	--------------

CODEINE SULFATE

SOLUTION;ORAL

CODEINE SULFATE

HIKMA	30MG/5ML	N202245 001	Jun 30, 2011
-------	----------	-------------	--------------

COLCHICINE

TABLET;ORAL

COLCRYS

+ TAKEDA PHARMS USA	0.6MG	N022352 001	Jul 29, 2009
---------------------	-------	-------------	--------------

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

COLCHICINE; PROBENECID

TABLET; ORAL

COLBENEMID

+ MERCK

0.5MG;500MG **

N012383 001

PROBEN-C

WATSON LABS

0.5MG;500MG

A085552 001

PROBENECID AND COLCHICINE

ANI PHARMS

0.5MG;500MG

A083734 001

BEECHAM

0.5MG;500MG

A084321 001

IMPAX LABS

0.5MG;500MG

A083720 002

SANDOZ

0.5MG;500MG

A086130 001

PROBENECID W/ COLCHICINE

LEDERLE

0.5MG;500MG

A086954 001

WATSON LABS

0.5MG;500MG

A083221 001

COLESEVELAM HYDROCHLORIDE

BAR, CHEWABLE; ORAL

WELCHOL

+ COSETTE

3.75GM

N210895 001 Apr 03, 2019

CAPSULE; ORAL

WELCHOL

COSETTE

375MG

N021141 001 May 26, 2000

FOR SUSPENSION; ORAL

COLESEVELAM HYDROCHLORIDE

IMPAX

3.75GM/PACKET

A212886 001 Jan 18, 2023

WATSON LABS INC

1.875GM/PACKET

A202178 001 Sep 01, 2020

3.75GM/PACKET

A202178 002 Sep 01, 2020

WELCHOL

+ COSETTE

1.875GM/PACKET **

N022362 001 Oct 02, 2009

TABLET; ORAL

COLESEVELAM HYDROCHLORIDE

CHARTWELL RX

625MG

A201354 001 Dec 17, 2020

INVAGEN PHARMS

625MG

A212602 001 Apr 20, 2020

UNITED RES LABS

625MG

A213456 001 Jan 21, 2022

WATSON LABS INC

625MG

A200830 001 Sep 02, 2020

COLESTIPOL HYDROCHLORIDE

GRANULE; ORAL

FLAVORED COLESTID

+ PFIZER

5GM/PACKET

N017563 001

+

5GM/SCOOPFUL

N017563 002

COLISTIN SULFATE

SUSPENSION; ORAL

COLY-MYCIN S

PARKE DAVIS

EQ 25MG BASE/5ML

N050355 001

CONIVAPTAN HYDROCHLORIDE

INJECTABLE; INTRAVENOUS

VAPRISOL

CUMBERLAND

20MG/4ML (5MG/ML)

N021697 001 Dec 29, 2005

COPANLISIB DIHYDROCHLORIDE

POWDER; INTRAVENOUS

ALIQOPA

+ BAYER HEALTHCARE

60MG/VIAL

N209936 001 Sep 14, 2017

COPPER

INTRAUTERINE DEVICE; INTRAUTERINE

CU-7

GD SEARLE LLC

89MG

N017408 001

TATUM-T

GD SEARLE LLC

120MG

N018205 001

CORTICOTROPIN

INJECTABLE; INJECTION

ACTH

PARKE DALE

25 UNITS/VIAL

N008317 002

40 UNITS/VIAL

N008317 004

ACTHAR

SANOFI AVENTIS US

25 UNITS/VIAL

N007504 002

40 UNITS/VIAL

N007504 003

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CORTICOTROPIN

INJECTABLE; INJECTION

ACTHAR GEL

MALLINCKRODT IRELAND 40 UNITS/ML N008372 006

CORTICOTROPIN

ORGANICS LAGRANGE 40 UNITS/ML N010831 001

80 UNITS/ML N010831 002

WATSON LABS 40 UNITS/VIAL A088772 001 Nov 21, 1984

PURIFIED CORTROPHIN GEL

ANI PHARMS 40 UNITS/ML N008975 001

CORTICOTROPIN-ZINC HYDROXIDE

INJECTABLE; INJECTION

CORTROPHIN-ZINC

ANI PHARMS 40 UNITS/ML N009854 001

CORTISONE ACETATE

INJECTABLE; INJECTION

CORTISONE ACETATE

PHARMACIA AND UPJOHN 25MG/ML N008126 002

WATSON LABS 25MG/ML A083147 003

25MG/ML A085677 001

50MG/ML A083147 004

50MG/ML A085677 002

CORTONE

MERCK 25MG/ML N007110 002

50MG/ML N007110 003

TABLET; ORAL

CORTISONE ACETATE

BARR 25MG A083471 001

CHARTWELL MOLECULAR 25MG A080694 001

ELKINS SINN 25MG A080836 001

EVERYLIFE 25MG A084246 001

HEATHER 25MG A085736 001

HIKMA INTL PHARMS 25MG A080776 002

IMPAX LABS 25MG N009458 001

INWOOD LABS 25MG A080731 001

IVAX SUB TEVA PHARMS 25MG A080630 001

25MG A083536 001

PANRAY 5MG N008284 002

25MG N008284 001

PHARMACIA AND UPJOHN 5MG N008126 003

10MG N008126 004

25MG N008126 001

PUREPAC PHARM 25MG A080493 001

VITARINE 25MG A080333 001

WATSON LABS 25MG A085884 001

WHITEWORTH TOWN PLSN 25MG A080341 001

CORTONE

+ MERCK 25MG ** N007750 003

COSYNTROPIN

INJECTABLE; INJECTION

COSYNTROPIN

NORVIUM BIOSCIENCE 0.25MG/VIAL A090574 001 Dec 17, 2009

SOLUTION; INTRAVENOUS

COSYNTROPIN

SANDOZ 0.25MG/ML (0.25MG/ML) N022028 001 Feb 21, 2008

CROMOLYN SODIUM

AEROSOL, METERED; INHALATION

INTAL

KING PHARMS LLC 0.8MG/INH N018887 001 Dec 05, 1985

CAPSULE; INHALATION

INTAL

+ SANOFI AVENTIS US 20MG ** N016990 001

CAPSULE; ORAL

GASTROCROM

UCB INC 100MG N019188 001 Dec 22, 1989

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CROMOLYN SODIUM

CONCENTRATE; ORAL

CROMOLYN SODIUM

GENERA PHARMS

10MG/5ML

A090954 001 Dec 18, 2009

SOLUTION; INHALATION

CROMOLYN SODIUM

ACTAVIS MID ATLANTIC

10MG/ML

A075067 001 Jul 19, 1999

BAUSCH

10MG/ML

A075585 001 Dec 21, 2000

EUGIA PHARMA

10MG/ML

A074209 001 Apr 26, 1994

HIKMA

10MG/ML

A075333 001 Apr 30, 2002

ROXANE

10MG/ML

A075175 001 Sep 30, 1999

WATSON LABS

10MG/ML

A076469 001 Jun 17, 2005

INTAL

+

KING PHARMS LLC

10MG/ML **

N018596 001 May 28, 1982

SOLUTION/DROPS; OPHTHALMIC

CROLOM

BAUSCH AND LOMB

4%

A074443 001 Jan 30, 1995

CROMOLYN SODIUM

APOTEX INC

4%

A075615 001 Jan 26, 2001

SCIEGEN PHARMS INC

4%

A074706 001 Apr 29, 1998

CROMOPTIC

KING PHARMS

4%

A075088 001 Apr 27, 1999

OPTICROM

+

ALLERGAN

4% **

N018155 001 Oct 03, 1984

SPRAY, METERED; NASAL

CROMOLYN SODIUM

ACTAVIS MID ATLANTIC

5.2MG/SPRAY

A074800 001 Jul 26, 2001

HH AND P

5.2MG/SPRAY

A077976 001 Sep 07, 2007

PERRIGO

5.2MG/SPRAY

A075427 001 Dec 12, 2001

NASALCROM

+

BLACKSMITH BRANDS

5.2MG/SPRAY **

N020463 001 Jan 03, 1997

CROTAMITON

CREAM; TOPICAL

EURAX

+

JOURNEY

10%

N006927 001

CRYPTENAMINE ACETATES

INJECTABLE; INJECTION

UNITENSEN

MEDPOINTE PHARM HLC

260CSR UNIT/ML

N008814 001

CRYPTENAMINE TANNATES

TABLET; ORAL

UNITENSEN

MEDPOINTE PHARM HLC

260CSR UNIT

N009217 001

CUPRIC SULFATE

INJECTABLE; INJECTION

CUPRIC SULFATE

+

ABRAXIS PHARM

EQ 0.4MG COPPER/ML **

N019350 001 May 05, 1987

CYANOCOBALAMIN

GEL, METERED; NASAL

NASCOBAL

ENDO OPERATIONS

0.5MG/INH

N019722 001 Nov 05, 1996

INJECTABLE; INJECTION

BERUBIGEN

PHARMACIA AND UPJOHN

1MG/ML

N006798 001

BETALIN 12

LILLY

0.1MG/ML

A080855 001

1MG/ML

A080855 002

COBAVITE

WATSON LABS

0.1MG/ML

A083013 001

1MG/ML

A083064 001

CYANOCOBALAMIN

ABRAXIS PHARM

0.03MG/ML

A080510 003

0.1MG/ML

A080510 001

1MG/ML

A080510 002

DELL LABS

0.03MG/ML

A080689 001

0.1MG/ML

A080689 002

1MG/ML

A080689 003

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CYANOCOBALAMIN

INJECTABLE; INJECTION

CYANOCOBALAMIN

DR REDDYS	0.1MG/ML	A080573 002	
	1MG/ML	A080573 001	
EPIC PHARMA LLC	1MG/ML	A087969 001	Nov 10, 1983
EUGIA PHARMA	1MG/ML	A213874 001	Dec 08, 2020
FRESENIUS KABI USA	0.1MG/ML	A080557 002	
LUITPOLD	0.03MG/ML	A080668 001	
LYPHOMED	1MG/ML	A083075 001	
MYLAN INSTITUTIONAL	1MG/ML	A040451 001	Sep 23, 2003
SANOFI AVENTIS US	1MG/ML	A080564 001	
SOLOPAK	1MG/ML	A087551 001	Feb 29, 1984
WARNER CHILCOTT	1MG/ML	N007085 002	
WATSON LABS	0.1MG/ML	A083120 001	
	1MG/ML	A083120 002	
WYETH AYERST	0.1MG/ML	A080554 001	
	1MG/ML	A080554 002	
XIROMED	1MG/ML	A215046 001	Aug 20, 2021
REDISOL			
MERCK	1MG/ML	N006668 010	
RUBIVITE			
BEL MAR	0.03MG/ML	N010791 004	
	0.05MG/ML	N010791 001	
	0.1MG/ML	N010791 002	
	0.12MG/ML	N010791 005	
	1MG/ML	N010791 003	
RUBRAMIN PC			
BRISTOL MYERS SQUIBB	0.1MG/ML	N006799 002	
+	1MG/ML **	N006799 004	
+	1MG/ML **	N006799 010	Apr 28, 1988
RUVITE			
SAVAGE LABS	1MG/ML	A080570 002	
VI-TWEL			
BAYER HLTHCARE	1MG/ML	N007012 002	
SPRAY, METERED; NASAL			
CALOMIST			
ENDO OPERATIONS	25MCG/SPRAY	N022102 001	Jul 27, 2007
NASCOBAL			
+	ENDO OPERATIONS	0.5MG/SPRAY	N021642 001
			Jan 31, 2005
TABLET; ORAL			
CYANOCOBALAMIN			
WEST WARD	1MG	A084264 001	

CYANOCOBALAMIN; CYANOCOBALAMIN CO-57; CYANOCOBALAMIN CO-58

N/A; N/A

DICOPAC KIT

GE HEALTHCARE	N/A; N/A; N/A	N017406 001	
---------------	---------------	-------------	--

CYANOCOBALAMIN; TANNIC ACID; ZINC ACETATE

INJECTABLE; INJECTION

DEPINAR

ARMOUR PHARM	0.5MG/ML; 2.3MG/ML; 1MG/ML	N011208 001	
--------------	----------------------------	-------------	--

CYCLACILLIN

FOR SUSPENSION; ORAL

CYCLAPEN-W

WYETH AYERST	125MG/5ML	N050508 001	
	250MG/5ML	N050508 002	
	500MG/5ML	N050508 003	

TABLET; ORAL

CYCLACILLIN

TEVA	250MG	A062895 001	Aug 04, 1988
	500MG	A062895 002	Aug 04, 1988

CYCLAPEN-W

WYETH AYERST	250MG	N050509 001	
	500MG	N050509 002	

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CYCLIZINE LACTATE

INJECTABLE; INJECTION

MAREZINE

GLAXOSMITHKLINE 50MG/ML N009495 001

CYCLOBENZAPRINE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

CYCLOBENZAPRINE HYDROCHLORIDE

APOTEX 15MG A206703 001 Jul 24, 2018
30MG A206703 002 Jul 24, 2018

TABLET; ORAL

CYCLOBENZAPRINE HYDROCHLORIDE

AIPING PHARM INC 5MG A072854 002 Feb 03, 2006
10MG A072854 001 Nov 19, 1991
PLIVA 10MG A074421 001 Sep 29, 1995
RISING 5MG A073144 002 Feb 03, 2006
7.5MG A073144 003 Mar 25, 2013
10MG A073144 001 May 30, 1991
SANDOZ 10MG A073683 001 Feb 26, 1993
WATSON LABS 10MG A073143 001 Nov 27, 1991
10MG A074436 001 Nov 30, 1994

FLEXERIL

+ JANSSEN RES AND DEV 5MG ** N017821 001
+ 10MG ** N017821 002CYCLOPENTOLATE HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

AK-PENTOLATE

EPIC PHARMA LLC 1% A085555 001

CYCLOPENTOLATE HYDROCHLORIDE

ALCON PHARMS LTD 1% A089162 001 Jan 24, 1991
SCIEGEN PHARMS INC 0.5% A205937 001 Dec 09, 2015
SOLA BARNES HIND 1% A084150 001
1% A084863 001

PENTOLAIR

PHARMAFAIR 0.5% A088643 001 Feb 09, 1987
1% A088150 001 Feb 25, 1983CYCLOPHOSPHAMIDE

CAPSULE; ORAL

CYCLOPHOSPHAMIDE

ANI PHARMS 25MG A207014 001 Mar 19, 2018
50MG A207014 002 Mar 19, 2018

INJECTABLE; INJECTION

CYCLOPHOSPHAMIDE

BAXTER HLTHCARE 100MG/VIAL A088371 001 Jul 03, 1986
200MG/VIAL A088372 001 Jul 03, 1986
500MG/VIAL A088373 001 Jul 03, 1986
1GM/VIAL A088374 001 Sep 24, 1986

CYTOXAN

+ BAXTER HLTHCARE 100MG/VIAL ** N012142 001
+ 200MG/VIAL ** N012142 002

CYTOXAN (LYOPHILIZED)

+ BAXTER HLTHCARE 500MG/VIAL ** N012142 003
+ 500MG/VIAL ** N012142 008 Jan 04, 1984
+ 1GM/VIAL ** N012142 004 Aug 30, 1982
+ 1GM/VIAL ** N012142 010 Sep 24, 1985
+ 2GM/VIAL ** N012142 005 Aug 30, 1982
+ 2GM/VIAL ** N012142 009 Dec 10, 1985

LYOPHILIZED CYTOXAN

+ BAXTER HLTHCARE 100MG/VIAL ** N012142 006 Dec 05, 1985
+ 200MG/VIAL ** N012142 007 Dec 10, 1985

NEOSAR

BEDFORD 100MG/VIAL A087442 001 Feb 16, 1982
200MG/VIAL A087442 002 Feb 16, 1982
500MG/VIAL A087442 003 Feb 16, 1982
1GM/VIAL A087442 004 Jul 08, 1983
2GM/VIAL A087442 005 Mar 30, 1989
TEVA PARENTERAL 100MG/VIAL A040015 001 Apr 29, 1993
200MG/VIAL A040015 002 Apr 29, 1993
500MG/VIAL A040015 003 Apr 29, 1993

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CYCLOPHOSPHAMIDE

INJECTABLE; INJECTION

NEOSAR

1GM/VIAL

A040015 004 Apr 29, 1993

2GM/VIAL

A040015 005 Apr 29, 1993

TABLET; ORAL

CYCLOPHOSPHAMIDE

ROXANE

25MG

A040032 001 Aug 17, 1999

50MG

A040032 002 Aug 17, 1999

CYCLOSPORINE

CAPSULE; ORAL

GENGRAF

ABBVIE

50MG

A065003 002 May 12, 2000

NEORAL

+ NOVARTIS

50MG **

N050715 003 Jul 14, 1995

SOLUTION; ORAL

CYCLOSPORINE

DR REDDYS LABS SA

100MG/ML

A065054 001 Dec 18, 2001

PHARM ASSOC

100MG/ML

A065167 001 Jan 05, 2005

PHARMOBEDIANT CNSLTG

100MG/ML

A065133 001 Sep 17, 2004

CYCLOTHIAZIDE

TABLET; ORAL

ANHYDRON

LILLY

2MG

N013157 002

FLUIDIL

PHARMACIA AND UPJOHN

2MG

N018173 001

CYCRIMINE HYDROCHLORIDE

TABLET; ORAL

PAGITANE

LILLY

1.25MG

N008951 001

2.5MG

N008951 002

CYPROHEPTADINE HYDROCHLORIDE

SYRUP; ORAL

CYPROHEPTADINE HYDROCHLORIDE

+ ACTAVIS MID ATLANTIC 2MG/5ML **

A086833 001

HALSEY

2MG/5ML

A089199 001 Jul 03, 1986

MORTON GROVE

2MG/5ML

A087001 001 Nov 04, 1982

NASKA

2MG/5ML

A089021 001 Dec 21, 1987

PATRIN

2MG/5ML

A204823 001 Dec 27, 2016

PHARM ASSOC

2MG/5ML

A091295 001 Mar 28, 2013

TRIS PHARMA INC

2MG/5ML

A205431 001 Dec 21, 2021

PERIACTIN

+ MERCK 2MG/5ML **

N013220 002

TABLET; ORAL

CYPROHEPTADINE HYDROCHLORIDE

AM THERAP

4MG

A088798 001 Feb 15, 1985

ASCOT

4MG

A087685 001 Oct 25, 1982

CHARTWELL RX

4MG

A086808 001

DURAMED PHARMS BARR

4MG

A088232 001 Oct 25, 1983

HALSEY

4MG

A089057 001 Jul 03, 1986

+ HERITAGE PHARMA

4MG

A087056 001

KV PHARM

4MG

A086737 001

MD PHARM

4MG

A087566 001 Nov 10, 1982

MYLAN

4MG

A086678 001

PIONEER PHARMS

4MG

A087839 001 Feb 08, 1984

PLIVA

4MG

A088205 001 Jul 26, 1983

RISING

4MG

A207783 001 Dec 29, 2016

STRIDES PHARMA

4MG

A087129 001

SUPERPHARM

4MG

A087405 001

VITARINE

4MG

A087284 001

WATSON LABS

4MG

A085245 001

4MG

A086165 001

4MG

A086580 001

PERIACTIN

+ MERCK 4MG **

N012649 001

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CYSTEINE HYDROCHLORIDE

INJECTABLE; INJECTION

CYSTEINE HYDROCHLORIDE

+ HOSPIRA 7.25% ** N019523 001 Oct 22, 1986

SOLUTION; INTRAVENOUS

ELCYS

+ EXELA PHARMA 2500MG/50ML (50MG/ML) N210660 002 Dec 04, 2023

NOURESS

+ BAXTER HLTHCARE CORP 500MG/10ML (50MG/ML) N212535 001 Dec 13, 2019

CYTARABINE

INJECTABLE; INJECTION

CYTARABINE

MEITHEAL 20MG/ML A206189 001 Jun 26, 2020

20MG/ML A206190 001 Nov 09, 2017

NORVIUM BIOSCIENCE 20MG/ML A200916 001 Dec 13, 2011

RISING 20MG/ML A200914 001 Dec 13, 2011

+ TEVA PARENTERAL 100MG/VIAL ** N016793 001

+ 500MG/VIAL ** N016793 002

+ 1GM/VIAL ** N016793 003 Dec 21, 1987

+ 2GM/VIAL ** N016793 004 Dec 21, 1987

CYTOSAR-U

TEVA PHARMS USA 100MG/VIAL A075206 001 Dec 30, 1998

500MG/VIAL A075206 002 Dec 30, 1998

1GM/VIAL A075206 004 Dec 30, 1998

2GM/VIAL A075206 003 Dec 30, 1998

INJECTABLE, LIPOSOMAL; INJECTION

DEPOCYT

+ PACIRA PHARMS INC 10MG/ML N021041 001 Apr 01, 1999

DACARBAZINE

INJECTABLE; INJECTION

DACARBAZINE

ABRAXIS PHARM 100MG/VIAL A070962 001 Aug 28, 1986

200MG/VIAL A070990 001 Aug 28, 1986

DTIC-DOME

+ BAYER HLTHCARE 100MG/VIAL ** N017575 001

+ 200MG/VIAL ** N017575 002

DACLATASVIR DIHYDROCHLORIDE

TABLET; ORAL

DAKLINZA

+ BRISTOL-MYERS SQUIBB EQ 30MG BASE N206843 001 Jul 24, 2015

+ EQ 60MG BASE N206843 002 Jul 24, 2015

+ EQ 90MG BASE N206843 003 Apr 13, 2016

DACTINOMYCIN

INJECTABLE; INJECTION

COSMEGEN

+ RECORDATI RARE 0.5MG/VIAL N050682 001

DACTINOMYCIN

AM REGENT 0.5MG/VIAL A202562 001 Aug 23, 2013

HIKMA 0.5MG/VIAL A090304 001 Mar 16, 2010

DALFAMPRIDINE

TABLET, EXTENDED RELEASE; ORAL

DALFAMPRIDINE

HIKMA 10MG A206646 001 Oct 24, 2018

RISING 10MG A206858 001 Jul 06, 2020

DALFOPRISTIN; QUINUPRISTIN

INJECTABLE; INTRAVENOUS

SYNERCID

+ KING PHARMS 350MG/VIAL; 150MG/VIAL N050748 001 Sep 21, 1999

420MG/VIAL; 180MG/VIAL N050748 002 Aug 24, 2000

DALTEPARIN SODIUM

INJECTABLE; INJECTION

FRAGMIN

PFIZER 7,500 IU/0.75ML N020287 008 Apr 04, 2002

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DALTEPARIN SODIUM

INJECTABLE;SUBCUTANEOUS

FRAGMIN

PFIZER	10,000IU/0.4ML (25,000IU/ML)	N020287 002	May 01, 2007
	95,000IU/9.5ML (10,000IU/ML)	N020287 007	Apr 04, 2002

DANAPAROID SODIUM

INJECTABLE;INJECTION

ORGARAN

ASPEN GLOBAL INC	750 UNITS/0.6ML	N020430 001	Dec 24, 1996
------------------	-----------------	-------------	--------------

DANAZOL

CAPSULE;ORAL

DANAZOL

AM THERAP	200MG	A071569 001	Dec 30, 1987
-----------	-------	-------------	--------------

DANOCRINE

SANOFI AVENTIS US	50MG **	N017557 003	
	100MG **	N017557 004	
	200MG **	N017557 002	

DANTROLENE SODIUM

INJECTABLE;INJECTION

DANTROLENE SODIUM

EUGIA PHARMA SPECLTS	20MG/VIAL	A205239 001	Feb 18, 2016
----------------------	-----------	-------------	--------------

DAPAGLIFLOZIN; METFORMIN HYDROCHLORIDE; SAXAGLIPTIN HYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL

QTERNMET XR

+ ASTRAZENECA AB	2.5MG;1GM;EQ 2.5MG BASE	N210874 001	May 02, 2019
	5MG;1GM;EQ 2.5MG BASE	N210874 002	May 02, 2019
	5MG;1GM;EQ 5MG BASE	N210874 003	May 02, 2019
	10MG;1GM;EQ 5MG BASE	N210874 004	May 02, 2019

DAPIPRAZOLE HYDROCHLORIDE

SOLUTION/DROPS;OPHTHALMIC

DAPIPRAZOLE HYDROCHLORIDE

BARADAINA LLC	0.5%	A204902 001	May 30, 2019
+ FERA PHARMS	0.5% **	N019849 001	Dec 31, 1990

DAPRODUSTAT

TABLET;ORAL

JESDUVROQ

+ GLAXOSMITHKLINE	1MG	N216951 001	Feb 01, 2023
	2MG	N216951 002	Feb 01, 2023
	4MG	N216951 003	Feb 01, 2023
	6MG	N216951 004	Feb 01, 2023
	8MG	N216951 005	Feb 01, 2023

DAPSONE

GEL;TOPICAL

DAPSONE

PADAGIS ISRAEL	5%	A215087 001	Oct 31, 2023
	7.5%	A212657 001	Nov 01, 2023

TABLET;ORAL

DAPSONE

ALVOGEN	25MG	A205429 001	Jan 07, 2016
	100MG	A205429 002	Jan 07, 2016
CHARTWELL RX	25MG	A204074 001	May 10, 2016
	100MG	A204074 002	May 10, 2016
TARO	25MG	A209430 001	Mar 01, 2019
	100MG	A209430 002	Mar 01, 2019

DAPTOMYCIN

POWDER;INTRAVENOUS

CUBICIN

CUBIST PHARMS LLC	250MG/VIAL **	N021572 001	Sep 12, 2003
	500MG/VIAL **	N021572 002	Sep 12, 2003

CUBICIN RF

+ CUBIST PHARMS LLC	500MG/VIAL **	N021572 003	Jul 06, 2016
---------------------	---------------	-------------	--------------

DAPTOMYCIN

HOSPIRA	500MG/VIAL	A202857 001	Sep 12, 2014
---------	------------	-------------	--------------

DAPZURA RT

+ BAXTER HLTHCARE CORP	500MG/VIAL **	N213645 001	Jan 25, 2022
------------------------	---------------	-------------	--------------

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DARIFENACIN HYDROBROMIDE

TABLET, EXTENDED RELEASE;ORAL

DARIFENACIN HYDROBROMIDE

ENDO OPERATIONS	EQ 7.5MG BASE	A091190 001	Mar 13, 2015
	EQ 15MG BASE	A091190 002	Mar 13, 2015
JUBILANT GENERICS	EQ 7.5MG BASE	A205550 001	Oct 12, 2016
	EQ 15MG BASE	A205550 002	Oct 12, 2016
XIROMED	EQ 7.5MG BASE	A209571 002	Oct 22, 2019
	EQ 15MG BASE	A209571 001	Oct 22, 2019
ENABLEX			
+ ABBVIE	EQ 7.5MG BASE **	N021513 001	Dec 22, 2004
+	EQ 15MG BASE **	N021513 002	Dec 22, 2004

DARUNAVIR

TABLET;ORAL

PREZISTA

+ JANSSEN PRODS	300MG **	N021976 001	Jun 23, 2006
+	400MG **	N021976 003	Oct 21, 2008

DASABUVIR SODIUM; OMBITASVIR, PARITAPREVIR, RITONAVIR

TABLET;ORAL

VIEKIRA PAK (COPACKAGED)

+ ABBVIE	EQ 250MG BASE;12.5MG, 75MG, 50MG	N206619 001	Dec 19, 2014
----------	----------------------------------	-------------	--------------

DASABUVIR SODIUM; OMBITASVIR; PARITAPREVIR; RITONAVIR

TABLET, EXTENDED RELEASE;ORAL

VIEKIRA XR

+ ABBVIE	EQ 200MG BASE;8.33MG;50MG;33.33MG **	N208624 001	Jul 22, 2016
----------	--------------------------------------	-------------	--------------

DAUNORUBICIN CITRATE

INJECTABLE, LIPOSOMAL;INJECTION

DAUNOXOME

GALEN (UK)	EQ 2MG BASE/ML	N050704 002	Apr 08, 1996
------------	----------------	-------------	--------------

DAUNORUBICIN HYDROCHLORIDE

INJECTABLE;INJECTION

CERUBIDINE

SANOFI AVENTIS US	EQ 20MG BASE/VIAL	A061876 001	
WYETH AYERST	EQ 20MG BASE/VIAL **	N050484 001	

DAUNORUBICIN HYDROCHLORIDE

HISUN PHARM HANGZHOU	EQ 20MG BASE/VIAL	A206195 001	Apr 25, 2019
TEVA PARENTERAL	EQ 20MG BASE/VIAL	A064212 001	Jun 23, 1998
	EQ 50MG BASE/VIAL	A064212 002	May 03, 1999

DECAMETHONIUM BROMIDE

INJECTABLE;INJECTION

SYNCURINE

GLAXOSMITHKLINE	1MG/ML	N006931 002	
-----------------	--------	-------------	--

DECITABINE

INJECTABLE;INTRAVENOUS

DACOGEN

+ OTSUKA	50MG/VIAL **	N021790 001	May 02, 2006
----------	--------------	-------------	--------------

DECITABINE

CIPLA	50MG/VIAL	A208601 001	Nov 16, 2017
-------	-----------	-------------	--------------

DEFERASIROX

GRANULE;ORAL

DEFERASIROX

AMNEAL	90MG	A214194 003	Aug 02, 2021
	180MG	A214194 001	Feb 09, 2021
	360MG	A214194 002	Feb 09, 2021
NORVIUM BIOSCIENCE	90MG	A213146 001	May 03, 2024
	180MG	A213146 002	May 03, 2024
	360MG	A213146 003	May 03, 2024

TABLET;ORAL

DEFERASIROX

CIPLA	90MG	A211852 001	Feb 11, 2020
	180MG	A211852 003	Jun 15, 2020
	360MG	A211852 002	Feb 11, 2020
STEVENS J	90MG	A210727 001	Dec 27, 2019
	180MG	A210727 003	Jun 15, 2020
	360MG	A210727 002	Dec 27, 2019

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DEFERASIROX

TABLET; ORAL

DEFERASIROX

TEVA PHARMS USA	90MG	A209223 001	Nov 25, 2019
	180MG	A209223 003	Apr 24, 2020
	360MG	A209223 002	Nov 25, 2019

TABLET, FOR SUSPENSION; ORAL

DEFERASIROX

NORVIUM BIOSCIENCE	125MG	A206585 001	May 17, 2024
	250MG	A206585 002	May 17, 2024
	500MG	A206585 003	May 17, 2024
ZYDUS PHARMS	125MG	A208882 001	May 05, 2020
	250MG	A208882 002	May 05, 2020
	500MG	A208882 003	May 05, 2020

DEFERIPRONE

SOLUTION; ORAL

FERRIPROX

+ CHIESI	80MG/ML	N208030 002	Apr 20, 2018
----------	---------	-------------	--------------

DEFEROXAMINE MESYLATE

INJECTABLE; INJECTION

DEFEROXAMINE MESYLATE

DR REDDYS	500MG/VIAL	A076806 001	Mar 31, 2006
	2GM/VIAL	A076806 002	Mar 31, 2006

DESFERAL

+ NOVARTIS	2GM/VIAL **	N016267 002	May 25, 2000
------------	-------------	-------------	--------------

DELAVIRDINE MESYLATE

TABLET; ORAL

RESCRIPTOR

+ VIIV HLTHCARE	100MG	N020705 001	Apr 04, 1997
+	200MG	N020705 002	Jul 14, 1999

DEMECARIUM BROMIDE

SOLUTION/DROPS; OPHTHALMIC

HUMORSOL

MERCK	0.125%	N011860 002	
	0.25%	N011860 001	

DEMECLOCYCLINE HYDROCHLORIDE

CAPSULE; ORAL

DECLOMYCIN

LEDERLE	150MG	N050262 001	
---------	-------	-------------	--

SYRUP; ORAL

DECLOMYCIN

LEDERLE	75MG/5ML	N050257 001	
---------	----------	-------------	--

TABLET; ORAL

DECLOMYCIN

COREPHARMA	75MG **	N050261 001	
	150MG **	N050261 002	
	300MG **	N050261 003	

DEMECLOCYCLINE HYDROCHLORIDE

BARR	150MG	A065171 001	Dec 13, 2004
	300MG	A065171 002	Dec 13, 2004
IMPAX LABS	150MG	A065094 001	Mar 22, 2004
	300MG	A065094 002	Mar 22, 2004

DESERPIDINE

TABLET; ORAL

HARMONYL

ABBVIE	0.1MG	N010796 001	
	0.25MG	N010796 002	

DESERPIDINE; HYDROCHLOROTHIAZIDE

TABLET; ORAL

ORETICYL 25

ABBVIE	0.125MG; 25MG	N012148 001	
--------	---------------	-------------	--

ORETICYL 50

ABBVIE	0.125MG; 50MG	N012148 003	
--------	---------------	-------------	--

ORETICYL FORTE

ABBVIE	0.25MG; 25MG	N012148 002	
--------	--------------	-------------	--

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DESERPIDINE; METHYLCLOTHIAZIDE

TABLET; ORAL

ENDURONYL

ABBOTT 0.25MG; 5MG

N012775 001

ENDURONYL FORTE

ABBOTT 0.5MG; 5MG

N012775 002

METHYLCLOTHIAZIDE AND DESERPIDINE

WATSON LABS 0.25MG; 5MG

A088486 001 Aug 10, 1984

0.5MG; 5MG

A088452 001 Aug 10, 1984

DESIPRAMINE HYDROCHLORIDE

CAPSULE; ORAL

PERTOFRANE

SANOFI AVENTIS US 25MG

N013621 001

50MG

N013621 002

TABLET; ORAL

DESIPRAMINE HYDROCHLORIDE

ANI PHARMS 10MG

A205153 001 Oct 28, 2016

25MG

A071803 002 Dec 08, 1987

25MG

A205153 002 Oct 28, 2016

50MG

A071803 003 Dec 08, 1987

50MG

A205153 003 Oct 28, 2016

75MG

A071803 004 Dec 08, 1987

75MG

A205153 004 Oct 28, 2016

100MG

A071803 001 May 29, 1997

100MG

A205153 005 Oct 28, 2016

150MG

A071803 005 May 29, 1997

150MG

A205153 006 Oct 28, 2016

25MG

A071864 001 Sep 09, 1987

50MG

A071865 001 Sep 09, 1987

75MG

A071866 001 Sep 09, 1987

100MG

A071867 001 Sep 09, 1987

DESLANOSIDE

INJECTABLE; INJECTION

CEDILANID-D

NOVARTIS 0.2MG/ML

N009282 002

DESLORATADINE

SOLUTION; ORAL

CLARINEX

+ MERCK SHARP DOHME 0.5MG/ML **

N021300 001 Sep 01, 2004

DESLORATADINE

TARO 0.5MG/ML

A202592 001 Jun 30, 2015

0.5MG/ML

A202936 001 May 26, 2016

TABLET; ORAL

DESLORATADINE

CHARTWELL RX 5MG

A078364 001 Dec 03, 2010

DASH PHARMS 5MG

A078351 001 Feb 10, 2012

PERRIGO 5MG

A078361 001 Dec 22, 2011

SUN PHARM INDS 5MG

A078359 001 Nov 16, 2010

TABLET, ORALLY DISINTEGRATING; ORAL

CLARINEX

+ ORGANON 2.5MG **

N021312 002 Jul 14, 2005

+ 5MG **

N021312 001 Jun 26, 2002

DESLORATADINE; PSEUDOEPHEDRINE SULFATE

TABLET, EXTENDED RELEASE; ORAL

CLARINEX D 24 HOUR

+ ORGANON 5MG; 240MG **

N021605 001 Mar 03, 2005

DESMOPRESSIN ACETATE

INJECTABLE; INJECTION

DDAVP

FERRING PHARMS INC 0.015MG/ML

N018938 002 Apr 25, 1995

DESMOPRESSIN ACETATE

AM REGENT 0.004MG/ML

A091374 001 Feb 14, 2019

BEDFORD 0.004MG/ML

A074575 001 Feb 18, 2000

HOSPIRA 0.004MG/ML

A075220 001 Aug 28, 2000

SUN PHARM INDS LTD 0.004MG/ML

A091280 001 Jan 25, 2013

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DESMOPRESSIN ACETATE

INJECTABLE; INJECTION

DESMOPRESSIN ACETATE PRESERVATIVE FREE

BEDFORD 0.004MG/ML

A074574 001 Feb 18, 2000

SOLUTION; NASAL

CONCENTRAID

FERRING 0.01%

N019776 001 Dec 26, 1990

DDAVP

+ FERRING PHARMS INC 0.01%

N017922 001

DESMOPRESSIN ACETATE

SUN PHARM INDS 0.01%

A077212 001 Apr 12, 2012

SPRAY, METERED; NASAL

DDAVP

+ FERRING PHARMS INC 0.01MG/SPRAY **

N017922 002 Feb 06, 1989

DDAVP (NEEDS NO REFRIGERATION)

+ FERRING PHARMS INC 0.01MG/SPRAY

N017922 003 Aug 07, 1996

DESMOPRESSIN ACETATE (NEEDS NO REFRIGERATION)

SUN PHARM 0.01MG/SPRAY

A078271 001 Dec 23, 2013

MINIRIN

+ FERRING 0.01MG/SPRAY

N021333 001 Sep 16, 2002

NOCTIVA

+ ACERUS PHARMS 0.00083MG/SPRAY

N201656 001 Mar 03, 2017

+ 0.00166MG/SPRAY

N201656 002 Mar 03, 2017

STIMATE

FERRING PHARMS INC 0.15MG/SPRAY

N020355 001 Mar 07, 1994

STIMATE (NEEDS NO REFRIGERATION)

+ FERRING PHARMS INC 0.15MG/SPRAY

N020355 002 Oct 24, 2007

TABLET; ORAL

DESMOPRESSIN ACETATE

ACTAVIS LABS FL INC 0.1MG

A076470 001 Jul 01, 2005

0.2MG

A076470 002 Jul 01, 2005

FERRING 0.1MG

N021795 001 May 08, 2008

0.2MG

N021795 002 May 08, 2008

IMPAX LABS INC 0.1MG

A077122 001 Jan 25, 2006

0.2MG

A077122 002 Jan 25, 2006

NATCO PHARMA USA 0.1MG

A200653 001 Jun 27, 2014

0.2MG

A200653 002 Jun 27, 2014

TABLET; SUBLINGUAL

NOCDURNA

+ FERRING PHARMS INC 0.0277MG

N022517 001 Jun 21, 2018

+ 0.0553MG

N022517 002 Jun 21, 2018

DESOGESTREL; ETHINYL ESTRADIOL

TABLET; ORAL-21

DESOGEN

ORGANON USA INC 0.15MG; 0.03MG

N020071 001 Dec 10, 1992

DESOGESTREL AND ETHINYL ESTRADIOL

DURAMED PHARMS BARR 0.15MG; 0.03MG

A075256 001 Aug 12, 1999

ORTHO-CEPT

+ JANSSEN PHARMS 0.15MG; 0.03MG **

N020301 001 Dec 14, 1992

TABLET; ORAL-28

DESOGEN

ORGANON USA INC 0.15MG; 0.03MG

N020071 002 Dec 10, 1992

DESOGESTREL AND ETHINYL ESTRADIOL

DR REDDYS LABS SA 0.15MG, N/A; 0.02MG, 0.01MG

A076916 001 Dec 29, 2008

0.1MG, 0.125MG, 0.15MG; 0.025MG, 0.025MG, 0.

A077182 001 Jan 24, 2006

0.025MG

XIROMED 0.15MG; 0.03MG

A202085 001 May 20, 2015

EMOQUETTE

ENDO OPERATIONS 0.15MG; 0.03MG

A076675 001 Feb 25, 2011

KIMIDESS

ENDO OPERATIONS 0.15MG, N/A; 0.02MG, 0.01MG

A076681 001 Apr 30, 2015

MIRCETTE

+ TEVA BRANDED PHARM 0.15MG, N/A; 0.02MG, 0.01MG **

N020713 001 Apr 22, 1998

ORTHO-CEPT

+ JANSSEN PHARMS 0.15MG; 0.03MG **

N020301 002 Dec 14, 1992

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DESONIDE

AEROSOL, FOAM;TOPICAL

VERDESO

+ ALMIRALL

0.05%

N021978 001 Sep 19, 2006

GEL;TOPICAL

DESONATE

+ LEO PHARMA AS

0.05% **

N021844 001 Oct 20, 2006

LOTION;TOPICAL

DESONIDE

PAI HOLDINGS PHARM

0.05%

A207855 001 Sep 28, 2017

OINTMENT;TOPICAL

DESONIDE

HIKMA

0.05%

A208836 001 Mar 27, 2017

PAI HOLDINGS PHARM

0.05%

A212002 001 Mar 12, 2019

DESOXIMETASONE

CREAM;TOPICAL

DESOXIMETASONE

COSETTE

0.25%

A209595 001 Mar 04, 2020

SCIEGEN PHARMS INC

0.05%

A203787 001 Jan 06, 2017

0.25%

A203234 001 Jun 12, 2015

TOPICORT

+ TARO

0.25% **

N017856 001

TOPICORT LP

+ TARO

0.05% **

N018309 001

GEL;TOPICAL

TOPICORT

+ TARO

0.05% **

N018586 001 Mar 29, 1982

OINTMENT;TOPICAL

DESOXIMETASONE

ALTANA

0.25%

A073440 001 Apr 01, 1998

COSETTE

0.25%

A206740 001 Dec 23, 2016

PAI HOLDINGS PHARM

0.25%

A208101 001 Feb 25, 2016

TOPICORT

TARO

0.25%

A074286 001 Jun 07, 1996

+

0.25% **

N018763 001 Sep 30, 1983

DESOXYCORTICOSTERONE ACETATE

INJECTABLE;INJECTION

DOCA

ORGANON USA INC

5MG/ML

N001104 001

PELLET;IMPLANTATION

PERCORTEN

NOVARTIS

125MG

N005151 001

DESOXYCORTICOSTERONE PIVALATE

INJECTABLE;INJECTION

PERCORTEN

NOVARTIS

25MG/ML

N008822 001

DESVENLAFAXINE

TABLET, EXTENDED RELEASE;ORAL

KHEDEZLA

OSMOTICA PHARM CORP

50MG

N204683 001 Jul 10, 2013

100MG

N204683 002 Jul 10, 2013

DESVENLAFAXINE FUMARATE

TABLET, EXTENDED RELEASE;ORAL

DESVENLAFAXINE

+ SUN PHARM

EQ 50MG BASE

N205583 001 Jan 28, 2014

+

TEVA PHARMS USA

EQ 100MG BASE

N205583 002 Jan 28, 2014

EQ 50MG BASE

N205208 001 Oct 11, 2013

EQ 100MG BASE

N205208 002 Oct 11, 2013

DESVENLAFAXINE SUCCINATE

TABLET, EXTENDED RELEASE;ORAL

DESVENLAFAXINE SUCCINATE

NORVIUM BIOSCIENCE

EQ 50MG BASE

A204095 001 Jun 29, 2015

EQ 100MG BASE

A204095 002 Jun 29, 2015

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DEXAMETHASONE

AEROSOL; TOPICAL

AEROSEB-DEX

ALLERGAN HERBERT 0.01% ** A083296 002

DECASPRAY

+ MERCK 0.04% ** N012731 002

ELIXIR; ORAL

DECADRON

MERCK 0.5MG/5ML N012376 002

DEXAMETHASONE

ALPHARMA US PHARMS 0.5MG/5ML A088997 001 Oct 10, 1986

+ PHARMOBEDIANT CNSLTG 0.5MG/5ML A088254 001 Jul 27, 1983

HEXADROL

ASPEN GLOBAL INC 0.5MG/5ML N012674 001

GEL; TOPICAL

DECADERM

MERCK 0.1% N013538 001

SUSPENSION/DROPS; OPHTHALMIC

DEXAMETHASONE

WATSON LABS 0.1% A089170 001 May 09, 1989

TABLET; ORAL

DECADRON

+ MERCK 0.25MG ** N011664 004

+ 0.5MG ** N011664 001

+ 0.75MG ** N011664 002

+ 1.5MG ** N011664 003

+ 4MG ** N011664 005

+ 6MG ** N011664 006 Jul 30, 1982

DEXAMETHASONE

BAUSCH 1.5MG A040700 001 Aug 15, 2008

CHARTWELL RX 1.5MG A085456 001

IMPAX LABS 0.75MG A085376 001

KEY THERAP 1.5MG A088237 001 Apr 28, 1983

PANGEA 0.25MG A088149 001 Apr 28, 1983

PHOENIX LABS NY 0.75MG A083806 001

PRASCO 0.75MG A080399 001

PVT FORM 0.75MG A083420 001

ROXANE 0.25MG A084614 001

SUN PHARM INDUSTRIES 0.25MG A084013 001

0.25MG A084764 001

0.5MG A084084 001

0.5MG A084766 001

0.75MG A084081 001

0.75MG A084765 001

1.5MG A084086 001

1.5MG A084763 001

UPSHER SMITH 0.75MG A087534 001

1.5MG A087533 001

WATSON LABS 0.25MG A085455 001

0.5MG A085458 001

0.75MG A080968 001

0.75MG A084457 001

0.75MG A085818 001

1.5MG A085840 001

WHITWORTH TOWN PLSN 0.75MG A084327 001

DEXONE 0.5

SOLVAY 0.5MG A084991 001

DEXONE 0.75

SOLVAY 0.75MG A084993 001

DEXONE 1.5

SOLVAY 1.5MG A084990 001

DEXONE 4

SOLVAY 4MG A084992 001

HEXADROL

ASPEN GLOBAL INC 0.5MG N012675 004

0.75MG N012675 007

1.5MG N012675 009

4MG N012675 010

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DEXAMETHASONE ACETATE

INJECTABLE; INJECTION

DECADRON-LA

+ MERCK

EQ 8MG BASE/ML **

N016675 001

DEXAMETHASONE ACETATE

WATSON LABS

EQ 8MG BASE/ML

A084315 001

WATSON LABS TEVA

EQ 16MG BASE/ML

A087711 001 May 24, 1982

DEXAMETHASONE SODIUM PHOSPHATE

AEROSOL; NASAL

DEXACORT

UCB INC

EQ 0.1MG PHOSPHATE/INH

N014242 001

AEROSOL, METERED; INHALATION

DEXACORT

UCB INC

EQ 0.1MG PHOSPHATE/INH

N013413 001

CREAM; TOPICAL

DECADRON

MERCK

EQ 0.1% PHOSPHATE

N011983 002

INJECTABLE; INJECTION

DECADRON

+ MERCK

EQ 4MG PHOSPHATE/ML **

N012071 002

+

EQ 24MG PHOSPHATE/ML **

N012071 004

DEXACEN-4

CENT PHARMS

EQ 4MG PHOSPHATE/ML

A084342 001

DEXAMETHASONE

ABRAXIS PHARM

EQ 4MG PHOSPHATE/ML

A088448 001 Jan 25, 1984

FRESENIUS KABI USA

EQ 10MG PHOSPHATE/ML

A088469 001 Jan 25, 1984

DEXAMETHASONE SODIUM PHOSPHATE

BEL MAR

EQ 4MG PHOSPHATE/ML

A084752 001

DELL LABS

EQ 4MG PHOSPHATE/ML

A083161 001

DR REDDYS

EQ 4MG PHOSPHATE/ML

A089169 001 Apr 09, 1986

EPIC PHARMA LLC

EQ 4MG PHOSPHATE/ML

A084493 001

GENEYORK PHARMS

EQ 4MG PHOSPHATE/ML

A214890 002 Jul 28, 2023

EQ 10MG PHOSPHATE/ML

A214890 001 Jul 07, 2023

HIKMA

EQ 4MG PHOSPHATE/ML

A211451 001 Aug 01, 2023

INTL MEDICATION

EQ 20MG PHOSPHATE/ML

A088522 001 Feb 17, 1984

+ LUITPOLD

EQ 4MG PHOSPHATE/ML

A087440 001 Jul 21, 1982

LYPHOMED

EQ 4MG PHOSPHATE/ML

A087065 001

TEVA PARENTERAL

EQ 4MG PHOSPHATE/ML

A081125 001 Aug 31, 1990

EQ 10MG PHOSPHATE/ML

A081126 001 Aug 31, 1990

WATSON LABS

EQ 4MG PHOSPHATE/ML

A083702 001

EQ 4MG PHOSPHATE/ML

A084355 001

EQ 10MG PHOSPHATE/ML

A087668 001 Jul 01, 1982

EQ 24MG PHOSPHATE/ML

A085606 001

WYETH AYERST

EQ 4MG PHOSPHATE/ML

A085641 001

HEXADROL

+ ASPEN GLOBAL INC

EQ 4MG PHOSPHATE/ML **

N014694 002

+

EQ 10MG PHOSPHATE/ML **

N014694 003

EQ 20MG PHOSPHATE/ML

N014694 004

OINTMENT; OPHTHALMIC

DECADRON

MERCK

EQ 0.05% PHOSPHATE

N011977 001

DEXAIR

PHARMAFAIR

EQ 0.05% PHOSPHATE

A088071 001 Dec 28, 1982

MAXIDEX

ALCON

EQ 0.05% PHOSPHATE

A083342 001

SOLUTION/DROPS; OPHTHALMIC

DEXAIR

PHARMAFAIR

EQ 0.1% PHOSPHATE

A088433 001 Dec 15, 1983

DEXAMETHASONE SODIUM PHOSPHATE

SOLA BARNES HIND

EQ 0.1% PHOSPHATE

A084170 001

EQ 0.1% PHOSPHATE

A084173 001

SOLUTION/DROPS; OPHTHALMIC, OTIC

DECADRON

+ MERCK

EQ 0.1% PHOSPHATE

N011984 001

DEXAMETHASONE SODIUM PHOSPHATE

SANDOZ

EQ 0.1% PHOSPHATE

A088771 001 Jan 16, 1985

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DEXAMETHASONE SODIUM PHOSPHATE

SOLUTION/DROPS;OTIC

DEXAMETHASONE SODIUM PHOSPHATE

EPIC PHARMA LLC EQ 0.1% PHOSPHATE A084855 001

DEXAMETHASONE SODIUM PHOSPHATE; LIDOCAINE HYDROCHLORIDE

INJECTABLE;INJECTION

DECADRON W/ XYLOCAINE

MERCCK EQ 4MG PHOSPHATE/ML;10MG/ML N013334 002

DEXAMETHASONE SODIUM PHOSPHATE; NEOMYCIN SULFATE

OINTMENT;OPHTHALMIC

NEODECADRON

MERCCK EQ 0.05% PHOSPHATE;EQ 3.5MG BASE/GM N050324 001

SOLUTION/DROPS;OPHTHALMIC

NEODECADRON

MERCCK EQ 0.1% PHOSPHATE;EQ 3.5MG BASE/ML N050322 001

NEOMYCIN SULFATE AND DEXAMETHASONE SODIUM PHOSPHATE

BAUSCH AND LOMB EQ 0.1% PHOSPHATE;EQ 3.5MG BASE/ML A064055 001 Oct 30, 1995

NEOMYCIN SULFATE-DEXAMETHASONE SODIUM PHOSPHATE

ALCON PHARMS LTD EQ 0.1% PHOSPHATE;EQ 3.5MG BASE/ML A062714 001 Jul 21, 1986

PHARMAFAIR EQ 0.1% PHOSPHATE;EQ 3.5MG BASE/ML A062539 001 Jan 10, 1985

DEXAMETHASONE; NEOMYCIN SULFATE; POLYMYXIN B SULFATE

OINTMENT;OPHTHALMIC

DEXACIDIN

NOVARTIS 0.1%;EQ 3.5MG BASE/GM;10,000 UNITS/GM A062566 001 Feb 22, 1985

DEXASPORIN

PHARMAFAIR 0.1%;EQ 3.5MG BASE/GM;10,000 UNITS/GM A062411 001 May 16, 1983

SUSPENSION/DROPS;OPHTHALMIC

DEXACIDIN

NOVARTIS 0.1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML A062544 001 Oct 29, 1984

DEXASPORIN

PHARMAFAIR 0.1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML A062428 001 May 18, 1983

NEOMYCIN AND POLYMYXIN B SULFATES AND DEXAMETHASONE

ALCON PHARMS LTD 0.1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML A062721 001 Nov 17, 1986

DEXAMETHASONE; TOBRAMYCIN

SUSPENSION/DROPS;OPHTHALMIC

TOBRAMYCIN AND DEXAMETHASONE

AMNEAL 0.1%;0.3% A212991 001 Jul 15, 2021

DEXBROMPHENIRAMINE MALEATE

SYRUP;ORAL

DISOMER

SCHERING 2MG/5ML N011814 002

TABLET;ORAL

DISOMER

SCHERING 2MG N011814 001

DEXBROMPHENIRAMINE MALEATE; PSEUDOEPHEDRINE SULFATE

TABLET;ORAL

DISOPHROL

SCHERING 2MG;60MG N012394 002

TABLET, EXTENDED RELEASE;ORAL

BROMPHERIL

COPLY PHARM 6MG;120MG A089116 001 Jan 22, 1987

DEXBROMPHENIRAMINE MALEATE AND PSEUDOEPHEDRINE SULFATE

AVANTHI INC 6MG;120MG A078648 001 Feb 27, 2013

DISOBROM

SANDOZ 6MG;120MG A070770 001 Sep 30, 1991

DISOPHROL

SCHERING PLOUGH 6MG;120MG N013483 004 Sep 13, 1982

DRIXORAL

+ SCHERING PLOUGH 6MG;120MG ** N013483 003 Sep 13, 1982

RESPORAL

PIONEER PHARMS 6MG;120MG A089139 001 Jun 16, 1988

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DEXCHLORPHENIRAMINE MALEATE

SYRUP;ORAL

DEXCHLORPHENIRAMINE MALEATE

+ PAI HOLDINGS PHARM 2MG/5ML

A088251 001 Mar 23, 1984

POLARAMINE

SCHERING 2MG/5ML

A086837 001 Jul 19, 1982

TABLET;ORAL

DEXCHLORPHENIRAMINE MALEATE

ANI PHARMS 2MG

A088682 001 Jan 17, 1986

POLARAMINE

+ SCHERING 2MG

A086835 001

DEXLANSOPRAZOLE

TABLET, ORALLY DISINTEGRATING, DELAYED RELEASE;ORAL

DEXILANT SOLUTAB

+ TAKEDA PHARMS USA 30MG

N208056 001 Jan 26, 2016

DEXMEDETOMIDINE HYDROCHLORIDE

INJECTABLE;INJECTION

DEXMEDETOMIDINE HYDROCHLORIDE

AM REGENT EQ 200MCG BASE/2ML (EQ 100MCG BASE/ML)

A203773 001 May 12, 2017

PIRAMAL CRITICAL EQ 200MCG BASE/2ML (EQ 100MCG BASE/ML)

A214794 001 Sep 03, 2021

WILSHIRE PHARMS INC EQ 80MCG BASE/20ML (EQ 4MCG BASE/ML)

A212791 003 May 08, 2020

DEXMETHYLPHENIDATE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE;ORAL

DEXMETHYLPHENIDATE HYDROCHLORIDE

AUROLIFE PHARMA LLC 5MG

A204266 001 Aug 25, 2015

10MG

A204266 002 Aug 25, 2015

15MG

A204266 003 Aug 25, 2015

20MG

A204266 004 Dec 21, 2015

25MG

A204266 005 May 04, 2021

30MG

A202580 001 Aug 28, 2013

35MG

A204266 006 May 04, 2021

40MG

A204266 007 Aug 25, 2015

TABLET;ORAL

DEXMETHYLPHENIDATE HYDROCHLORIDE

BIONPHARMA 5MG

A209754 001 Mar 24, 2020

10MG

A209754 002 Mar 24, 2020

LANNETT CO INC 2.5MG

A209468 001 Sep 25, 2017

5MG

A209468 002 Sep 25, 2017

10MG

A209468 003 Sep 25, 2017

TEVA PHARMS 2.5MG

A077107 003 Jan 29, 2007

5MG

A077107 001 Jan 29, 2007

10MG

A077107 002 Jan 29, 2007

DEXRAZOXANE HYDROCHLORIDE

INJECTABLE;INJECTION

TOTECT

+ CLINIGEN EQ 500MG BASE/VIAL **

N022025 001 Sep 06, 2007

ZINECARD

+ PFIZER EQ 250MG BASE/VIAL

N020212 001 May 26, 1995

+ EQ 500MG BASE/VIAL

N020212 002 May 26, 1995

DEXTROAMPHETAMINE SULFATE

CAPSULE;ORAL

DEXAMPEX

TEVA 15MG

A085355 001

CAPSULE, EXTENDED RELEASE;ORAL

DEXTROAMPHETAMINE SULFATE

ABLE 5MG

A076814 001 Aug 25, 2004

10MG

A076814 002 Aug 25, 2004

15MG

A076814 003 Aug 25, 2004

DR REDDYS LABS SA 5MG

A076137 001 Jan 18, 2002

10MG

A076137 002 Jan 18, 2002

15MG

A076137 003 Jan 18, 2002

NESHER PHARMS 5MG

A209111 001 Jun 27, 2017

10MG

A209111 002 Jun 27, 2017

15MG

A209111 003 Jun 27, 2017

NORVIUM BIOSCIENCE 5MG

A206735 001 Jan 27, 2016

10MG

A206735 002 Jan 27, 2016

15MG

A206735 003 Jan 27, 2016

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DEXTROAMPHETAMINE SULFATE

CAPSULE, EXTENDED RELEASE;ORAL

DEXTROAMPHETAMINE SULFATE

STRIDES PHARMA	5MG	A205077 001	Jun 21, 2019
	10MG	A205077 002	Jun 21, 2019
	15MG	A205077 003	Jun 21, 2019

ELIXIR;ORAL

DEXEDRINE

GLAXOSMITHKLINE	5MG/5ML **	A083902 001	
-----------------	------------	-------------	--

TABLET;ORAL

DEXAMPEX

TEVA	5MG	A083735 001	
	10MG	A083735 002	

DEXEDRINE

+ GLAXOSMITHKLINE	5MG **	A084935 001	
-------------------	--------	-------------	--

DEXTROAMPHETAMINE SULFATE

ANI PHARMS	5MG	A085370 001	
EPIC PHARMA LLC	5MG	A090652 001	Mar 07, 2014
	10MG	A090652 002	Mar 07, 2014
HALSEY	10MG	A083930 001	
LANNETT	5MG	A083903 001	
	10MG	A083903 003	
	15MG	A085652 001	
MAST MM	5MG	A086521 001	
NESHER PHARMS	5MG	A040365 001	Oct 31, 2002
	5MG	A206588 001	Mar 28, 2016
	10MG	A040367 001	Oct 31, 2002
	10MG	A206588 002	Mar 28, 2016
PUREPAC PHARM	5MG	A084125 001	
SANDOZ	10MG	A085371 001	
STRIDES PHARMA	5MG	A040299 001	May 13, 1999
TRIS PHARMA INC	5MG	A206095 001	Aug 18, 2022
	10MG	A206095 002	Aug 18, 2022
VITARINE	5MG	A084986 001	
	10MG	A085892 001	

DEXTROSTAT

+ SHIRE	5MG **	A084051 001	
---------	--------	-------------	--

+	10MG **	A084051 002	
---	---------	-------------	--

FERNDEX

FERNDALE LABS	5MG	A084001 001	
---------------	-----	-------------	--

DEXTROMETHORPHAN HYDROBROMIDE; PROMETHAZINE HYDROCHLORIDE

SYRUP;ORAL

PHERAZINE DM

HALSEY	15MG/5ML; 6.25MG/5ML	A088913 001	Mar 02, 1987
--------	----------------------	-------------	--------------

PROMETH W/ DEXTROMETHORPHAN

G AND W LABS INC	15MG/5ML; 6.25MG/5ML **	A088762 001	Oct 31, 1984
------------------	-------------------------	-------------	--------------

PROMETHAZINE HYDROCHLORIDE AND DEXTROMETHORPHAN HYDROBROMIDE

HIKMA	15MG/5ML; 6.25MG/5ML	A040027 001	Jul 31, 1996
-------	----------------------	-------------	--------------

DEXTROSE

INJECTABLE; INJECTION

DEXTROSE 10% IN PLASTIC CONTAINER

B BRAUN	10GM/100ML	N018046 001	
---------	------------	-------------	--

MILES	10GM/100ML	N018504 001	
-------	------------	-------------	--

DEXTROSE 2.5% IN PLASTIC CONTAINER

B BRAUN	2.5GM/100ML	N018358 001	
---------	-------------	-------------	--

	2.5GM/100ML	N019626 001	Feb 02, 1988
--	-------------	-------------	--------------

DEXTROSE 20% IN PLASTIC CONTAINER

+ BAXTER HLTHCARE	20GM/100ML **	N017521 004	
-------------------	---------------	-------------	--

DEXTROSE 30% IN PLASTIC CONTAINER

+ BAXTER HLTHCARE	30GM/100ML **	N017521 003	
-------------------	---------------	-------------	--

DEXTROSE 38.5% IN PLASTIC CONTAINER

ABBOTT	38.5GM/100ML	N018923 001	Sep 19, 1984
--------	--------------	-------------	--------------

DEXTROSE 40% IN PLASTIC CONTAINER

+ BAXTER HLTHCARE	40GM/100ML **	N017521 002	
-------------------	---------------	-------------	--

DEXTROSE 5%

AM REGENT	50MG/ML	A217263 001	Apr 08, 2024
-----------	---------	-------------	--------------

DEXTROSE 5% IN PLASTIC CONTAINER

DHL	5GM/100ML	N019971 001	Sep 28, 1995
-----	-----------	-------------	--------------

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DEXTROSE

INJECTABLE; INJECTION

DEXTROSE 5% IN PLASTIC CONTAINER			
+	ICU MEDICAL INC	50MG/ML	N019222 001 Jul 13, 1984
DEXTROSE 50% IN PLASTIC CONTAINER			
+	BAXTER HLTHCARE	50GM/100ML **	N017521 001
	ICU MEDICAL INC	50GM/100ML	N019894 001 Dec 26, 1989
DEXTROSE 60%			
	B BRAUN	60GM/100ML	N017995 002 Sep 22, 1982
DEXTROSE 60% IN PLASTIC CONTAINER			
	B BRAUN	60GM/100ML	N017995 001
+	BAXTER HLTHCARE	60GM/100ML **	N017521 005 Mar 26, 1982
		60GM/100ML	N020047 002 Jul 02, 1991
	HOSPIRA	60GM/100ML	N019346 001 Jan 25, 1985
DEXTROSE 7.7% IN PLASTIC CONTAINER			
	B BRAUN	7.7GM/100ML	N019626 003 Feb 02, 1988
DEXTROSE 70% IN PLASTIC CONTAINER			
+	BAXTER HLTHCARE	70GM/100ML **	N017521 006 Mar 26, 1982

DEXTROSE; MAGNESIUM ACETATE TETRAHYDRATE; POTASSIUM ACETATE; SODIUM CHLORIDE

INJECTABLE; INJECTION

PLASMA-LYTE 56 AND DEXTROSE 5% IN PLASTIC CONTAINER			
	BAXTER HLTHCARE	5GM/100ML; 32MG/100ML; 128MG/100ML; 234MG/100ML	N017385 001

DEXTROSE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; POTASSIUM PHOSPHATE, DIBASIC; SODIUM ACETATE

INJECTABLE; INJECTION

ISOLYTE P W/ DEXTROSE 5% IN PLASTIC CONTAINER			
	B BRAUN	5GM/100ML; 31MG/100ML; 130MG/100ML; 26MG/100ML; 320MG/100ML	N019025 001 Dec 27, 1984

DEXTROSE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; POTASSIUM PHOSPHATE, DIBASIC; SODIUM CHLORIDE; SODIUM LACTATE; SODIUM PHOSPHATE, MONOBASIC ANHYDROUS

INJECTABLE; INJECTION

IONOSOL B AND DEXTROSE 5% IN PLASTIC CONTAINER			
	HOSPIRA	5GM/100ML; 53MG/100ML; 100MG/100ML; 100MG/100ML; 180MG/100ML; 280MG/100ML; 16MG/100ML	N019515 001 May 08, 1986

DEXTROSE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE

INJECTABLE; INJECTION

ISOLYTE H IN DEXTROSE 5% IN PLASTIC CONTAINER			
	B BRAUN	5GM/100ML; 30MG/100ML; 97MG/100ML; 220MG/100ML; 140MG/100ML	N019844 001 Jun 10, 1993
ISOLYTE H W/ DEXTROSE 5% IN PLASTIC CONTAINER			
	B BRAUN	5GM/100ML; 30MG/100ML; 97MG/100ML; 220MG/100ML; 140MG/100ML	N018273 001

DEXTROSE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE; SODIUM GLUCONATE

INJECTABLE; INJECTION

ISOLYTE S IN DEXTROSE 5% IN PLASTIC CONTAINER			
	B BRAUN	5GM/100ML; 30MG/100ML; 37MG/100ML; 370MG/100ML; 530MG/100ML; 500MG/100ML	N019843 001 Aug 09, 1993
ISOLYTE S W/ DEXTROSE 5% IN PLASTIC CONTAINER			
	B BRAUN	5GM/100ML; 30MG/100ML; 37MG/100ML; 370MG/100ML; 530MG/100ML; 500MG/100ML	N018274 001
PLASMA-LYTE 148 AND DEXTROSE 5% IN PLASTIC CONTAINER			
+	BAXTER HLTHCARE	5GM/100ML; 30MG/100ML; 37MG/100ML; 368MG/100ML; 526MG/100ML; 502MG/100ML	N017451 001

DEXTROSE; POTASSIUM CHLORIDE

INJECTABLE; INJECTION

POTASSIUM CHLORIDE 0.037% IN DEXTROSE 5% IN PLASTIC CONTAINER			
	B BRAUN	5GM/100ML; 37MG/100ML	N019699 001 Sep 29, 1989
POTASSIUM CHLORIDE 0.075% IN DEXTROSE 5% IN PLASTIC CONTAINER			
	B BRAUN	5GM/100ML; 75MG/100ML	N018744 001 Nov 09, 1982
		5GM/100ML; 75MG/100ML	N019699 002 Sep 29, 1989
POTASSIUM CHLORIDE 0.11% IN DEXTROSE 5% IN PLASTIC CONTAINER			
	B BRAUN	5GM/100ML; 110MG/100ML	N019699 003 Sep 29, 1989
POTASSIUM CHLORIDE 0.15% IN DEXTROSE 5% IN PLASTIC CONTAINER			
	B BRAUN	5GM/100ML; 150MG/100ML	N018744 002 Nov 09, 1982
POTASSIUM CHLORIDE 0.22% IN DEXTROSE 5% IN PLASTIC CONTAINER			
	B BRAUN	5GM/100ML; 220MG/100ML	N018744 003 Nov 09, 1982

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DEXTROSE; POTASSIUM CHLORIDE

INJECTABLE; INJECTION

POTASSIUM CHLORIDE 0.22% IN DEXTROSE 5% IN PLASTIC CONTAINER			
5GM/100ML; 220MG/100ML		N019699 005	Sep 29, 1989
POTASSIUM CHLORIDE 0.3% IN DEXTROSE 5% IN PLASTIC CONTAINER			
B BRAUN	5GM/100ML; 300MG/100ML	N018744 004	Nov 09, 1982
POTASSIUM CHLORIDE 30MEQ IN DEXTROSE 5% IN PLASTIC CONTAINER			
ICU MEDICAL INC	5GM/100ML; 224MG/100ML	N018371 003	
POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% IN PLASTIC CONTAINER			
ICU MEDICAL INC	5GM/100ML; 298MG/100ML	N018371 002	

DEXTROSE; POTASSIUM CHLORIDE; POTASSIUM LACTATE; SODIUM CHLORIDE; SODIUM PHOSPHATE, MONOBASIC ANHYDROUS

INJECTABLE; INJECTION

IONOSOL T AND DEXTROSE 5% IN PLASTIC CONTAINER			
HOSPIRA	5GM/100ML; 111MG/100ML; 256MG/100ML; 146MG /100ML; 207MG/100ML	N019514 001	May 08, 1986

DEXTROSE; POTASSIUM CHLORIDE; POTASSIUM PHOSPHATE, DIBASIC; SODIUM ACETATE; SODIUM CHLORIDE

INJECTABLE; INJECTION

ISOLYTE M IN DEXTROSE 5% IN PLASTIC CONTAINER			
B BRAUN	5GM/100ML; 150MG/100ML; 130MG/100ML; 280MG /100ML; 91MG/100ML	N019870 001	Jun 10, 1993
ISOLYTE M W/ DEXTROSE 5% IN PLASTIC CONTAINER			
B BRAUN	5GM/100ML; 150MG/100ML; 130MG/100ML; 280MG /100ML; 91MG/100ML	N018270 001	

DEXTROSE; POTASSIUM CHLORIDE; POTASSIUM PHOSPHATE, MONOBASIC; SODIUM CHLORIDE; SODIUM LACTATE

INJECTABLE; INJECTION

DEXTROSE 5% AND ELECTROLYTE NO. 75 IN PLASTIC CONTAINER			
BAXTER HLTHCARE	5GM/100ML; 205MG/100ML; 100MG/100ML; 120MG /100ML; 220MG/100ML	N018840 001	Jun 29, 1983

DEXTROSE; POTASSIUM CHLORIDE; SODIUM CHLORIDE

INJECTABLE; INJECTION

DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 0.075%			
B BRAUN	5GM/100ML; 75MG/100ML; 200MG/100ML	N018268 009	
DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 0.15% IN PLASTIC CONTAINER			
B BRAUN	5GM/100ML; 150MG/100ML; 200MG/100ML	N018268 004	
DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 0.224% IN PLASTIC CONTAINER			
B BRAUN	5GM/100ML; 220MG/100ML; 200MG/100ML	N018268 005	
DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 0.3% IN PLASTIC CONTAINER			
B BRAUN	5GM/100ML; 300MG/100ML; 200MG/100ML	N018268 006	
DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 0.075% IN PLASTIC CONTAINER			
B BRAUN	5GM/100ML; 75MG/100ML; 330MG/100ML	N018268 011	Jan 18, 1986
DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 0.15% IN PLASTIC CONTAINER			
B BRAUN	5GM/100ML; 150MG/100ML; 330MG/100ML	N018268 012	Jan 18, 1986
DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 0.22% IN PLASTIC CONTAINER			
B BRAUN	5GM/100ML; 220MG/100ML; 330MG/100ML	N018268 013	Jan 18, 1986
DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 0.30% IN PLASTIC CONTAINER			
B BRAUN	5GM/100ML; 300MG/100ML; 330MG/100ML	N018268 014	Jan 18, 1986
DEXTROSE 5%, SODIUM CHLORIDE 0.45% AND POTASSIUM CHLORIDE 0.075%			
B BRAUN	5GM/100ML; 75MG/100ML; 450MG/100ML	N018268 010	
DEXTROSE 5%, SODIUM CHLORIDE 0.45% AND POTASSIUM CHLORIDE 0.15% IN PLASTIC CONTAINER			
B BRAUN	5GM/100ML; 150MG/100ML; 450MG/100ML	N018268 001	
DEXTROSE 5%, SODIUM CHLORIDE 0.45% AND POTASSIUM CHLORIDE 0.22% IN PLASTIC CONTAINER			
B BRAUN	5GM/100ML; 220MG/100ML; 450MG/100ML	N018268 002	
DEXTROSE 5%, SODIUM CHLORIDE 0.45% AND POTASSIUM CHLORIDE 0.3% IN PLASTIC CONTAINER			
B BRAUN	5GM/100ML; 300MG/100ML; 450MG/100ML	N018268 003	
DEXTROSE 5%, SODIUM CHLORIDE 0.45% AND POTASSIUM CHLORIDE 15MEQ IN PLASTIC CONTAINER			
BAXTER HLTHCARE	5GM/100ML; 224MG/100ML; 450MG/100ML	N018008 003	
DEXTROSE 5%, SODIUM CHLORIDE 0.45% AND POTASSIUM CHLORIDE 20MEQ (K) IN PLASTIC CONTAINER			
BAXTER HLTHCARE	5GM/100ML; 300MG/100ML; 450MG/100ML	N018008 001	
DEXTROSE 5%, SODIUM CHLORIDE 0.45% AND POTASSIUM CHLORIDE 5MEQ IN PLASTIC CONTAINER			
BAXTER HLTHCARE	5GM/100ML; 75MG/100ML; 450MG/100ML	N018008 002	
POTASSIUM CHLORIDE 10MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.225% IN PLASTIC CONTAINER			
+ ICU MEDICAL INC	5GM/100ML; 74.5MG/100ML; 225MG/100ML **	N018365 002	Jul 05, 1983
+ ICU MEDICAL INC	5GM/100ML; 149MG/100ML; 225MG/100ML **	N018365 006	Mar 28, 1988
POTASSIUM CHLORIDE 10MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER			
ICU MEDICAL INC	5GM/100ML; 74.5MG/100ML; 300MG/100ML	N018876 001	Jan 17, 1986
ICU MEDICAL INC	5GM/100ML; 149MG/100ML; 300MG/100ML	N018876 006	Mar 28, 1988

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DEXTROSE; POTASSIUM CHLORIDE; SODIUM CHLORIDE

INJECTABLE; INJECTION

POTASSIUM CHLORIDE 10MEQ	IN	DEXTROSE 5% AND SODIUM CHLORIDE 0.9%	IN	PLASTIC CONTAINER	
+	ICU MEDICAL INC	5GM/100ML;74.5MG/100ML;900MG/100ML			N019691 002 Mar 24, 1988
+		5GM/100ML;149MG/100ML;900MG/100ML			N019691 004 Mar 24, 1988
POTASSIUM CHLORIDE 15MEQ	IN	DEXTROSE 5% AND SODIUM CHLORIDE 0.225%	IN	PLASTIC CONTAINER	
	ICU MEDICAL INC	5GM/100ML;224MG/100ML;225MG/100ML **			N018365 008 Mar 28, 1988
POTASSIUM CHLORIDE 15MEQ	IN	DEXTROSE 5% AND SODIUM CHLORIDE 0.3%	IN	PLASTIC CONTAINER	
	ICU MEDICAL INC	5GM/100ML;224MG/100ML;300MG/100ML			N018876 007 Mar 28, 1988
POTASSIUM CHLORIDE 15MEQ	IN	DEXTROSE 5% AND SODIUM CHLORIDE 0.45%	IN	PLASTIC CONTAINER	
+	ICU MEDICAL INC	5GM/100ML;224MG/100ML;450MG/100ML			N018362 006 Mar 28, 1988
POTASSIUM CHLORIDE 15MEQ	IN	DEXTROSE 5% AND SODIUM CHLORIDE 0.9%	IN	PLASTIC CONTAINER	
+	ICU MEDICAL INC	5GM/100ML;224MG/100ML;900MG/100ML			N019691 006 Mar 24, 1988
POTASSIUM CHLORIDE 20MEQ	IN	DEXTROSE 5% AND SODIUM CHLORIDE 0.225%	IN	PLASTIC CONTAINER	
+	ICU MEDICAL INC	5GM/100ML;298MG/100ML;225MG/100ML **			N018365 009 Mar 28, 1988
POTASSIUM CHLORIDE 20MEQ	IN	DEXTROSE 5% AND SODIUM CHLORIDE 0.3%	IN	PLASTIC CONTAINER	
	ICU MEDICAL INC	5GM/100ML;298MG/100ML;300MG/100ML			N018876 008 Mar 28, 1988
POTASSIUM CHLORIDE 20MEQ	IN	DEXTROSE 5% AND SODIUM CHLORIDE 0.45%	IN	PLASTIC CONTAINER	
+	ICU MEDICAL INC	5GM/100ML;298MG/100ML;450MG/100ML			N018362 007 Mar 28, 1988
POTASSIUM CHLORIDE 20MEQ	IN	DEXTROSE 5% AND SODIUM CHLORIDE 0.9%	IN	PLASTIC CONTAINER	
+	ICU MEDICAL INC	5GM/100ML;298MG/100ML;900MG/100ML			N019691 008 Mar 24, 1988
POTASSIUM CHLORIDE 20MEQ	IN	DEXTROSE 5% IN SODIUM CHLORIDE 0.3%	IN	PLASTIC CONTAINER	
	ICU MEDICAL INC	5GM/100ML;149MG/100ML;300MG/100ML			N018876 002 Jan 17, 1986
POTASSIUM CHLORIDE 30MEQ	IN	DEXTROSE 5% AND SODIUM CHLORIDE 0.225%	IN	PLASTIC CONTAINER	
	ICU MEDICAL INC	5GM/100ML;224MG/100ML;225MG/100ML **			N018365 003 Jul 05, 1983
POTASSIUM CHLORIDE 30MEQ	IN	DEXTROSE 5% AND SODIUM CHLORIDE 0.3%	IN	PLASTIC CONTAINER	
	ICU MEDICAL INC	5GM/100ML;224MG/100ML;300MG/100ML			N018876 003 Jan 17, 1986
POTASSIUM CHLORIDE 30MEQ	IN	DEXTROSE 5% AND SODIUM CHLORIDE 0.9%	IN	PLASTIC CONTAINER	
+	ICU MEDICAL INC	5GM/100ML;224MG/100ML;900MG/100ML			N019691 007 Mar 24, 1988
POTASSIUM CHLORIDE 40MEQ	IN	DEXTROSE 5% AND SODIUM CHLORIDE 0.225%	IN	PLASTIC CONTAINER	
	ICU MEDICAL INC	5GM/100ML;298MG/100ML;225MG/100ML **			N018365 004 Jul 05, 1983
POTASSIUM CHLORIDE 40MEQ	IN	DEXTROSE 5% AND SODIUM CHLORIDE 0.3%	IN	PLASTIC CONTAINER	
	ICU MEDICAL INC	5GM/100ML;298MG/100ML;300MG/100ML			N018876 004 Mar 28, 1988
POTASSIUM CHLORIDE 5MEQ	IN	DEXTROSE 5% AND SODIUM CHLORIDE 0.225%	IN	PLASTIC CONTAINER	
	ICU MEDICAL INC	5GM/100ML;74.5MG/100ML;225MG/100ML **			N018365 005 Mar 28, 1988
		5GM/100ML;149MG/100ML;225MG/100ML **			N018365 007 Mar 28, 1988
POTASSIUM CHLORIDE 5MEQ	IN	DEXTROSE 5% AND SODIUM CHLORIDE 0.3%	IN	PLASTIC CONTAINER	
	ICU MEDICAL INC	5GM/100ML;74.5MG/100ML;300MG/100ML			N018876 005 Mar 28, 1988
		5GM/100ML;149MG/100ML;300MG/100ML			N018876 009 Mar 28, 1988
POTASSIUM CHLORIDE 5MEQ	IN	DEXTROSE 5% AND SODIUM CHLORIDE 0.45%	IN	PLASTIC CONTAINER	
+	ICU MEDICAL INC	5GM/100ML;74.5MG/100ML;450MG/100ML			N018362 008 Mar 28, 1988
+		5GM/100ML;149MG/100ML;450MG/100ML			N018362 004 Mar 28, 1988
POTASSIUM CHLORIDE 5MEQ	IN	DEXTROSE 5% AND SODIUM CHLORIDE 0.9%	IN	PLASTIC CONTAINER	
+	ICU MEDICAL INC	5GM/100ML;74.5MG/100ML;900MG/100ML			N019691 001 Mar 24, 1988
+		5GM/100ML;149MG/100ML;900MG/100ML			N019691 003 Mar 24, 1988

DEXTROSE; SODIUM CHLORIDE

INJECTABLE; INJECTION

DEXTROSE 10% AND SODIUM CHLORIDE 0.2%	IN	PLASTIC CONTAINER	
B BRAUN	10GM/100ML;200MG/100ML		N018386 001
DEXTROSE 10% AND SODIUM CHLORIDE 0.45%	IN	PLASTIC CONTAINER	
B BRAUN	10GM/100ML;450MG/100ML		N018229 001
DEXTROSE 10% AND SODIUM CHLORIDE 0.9%	IN	PLASTIC CONTAINER	
B BRAUN	10GM/100ML;900MG/100ML		N018047 001
BAXTER HLTHCARE	10GM/100ML;900MG/100ML		N016696 001
DEXTROSE 2.5% AND SODIUM CHLORIDE 0.45%	IN	PLASTIC CONTAINER	
B BRAUN	2.5GM/100ML;450MG/100ML		N018030 001
HOSPIRA	2.5GM/100ML;450MG/100ML		N018096 001
DEXTROSE 2.5% AND SODIUM CHLORIDE 0.9%	IN	PLASTIC CONTAINER	
B BRAUN	2.5GM/100ML;900MG/100ML		N018376 001
DEXTROSE 3.3% AND SODIUM CHLORIDE 0.3%	IN	PLASTIC CONTAINER	
ABBOTT	3.3GM/100ML;300MG/100ML		N018055 001
DEXTROSE 5% AND SODIUM CHLORIDE 0.11%	IN	PLASTIC CONTAINER	
B BRAUN	5GM/100ML;110MG/100ML		N018030 005
DEXTROSE 5% AND SODIUM CHLORIDE 0.2%	IN	PLASTIC CONTAINER	
B BRAUN	5GM/100ML;200MG/100ML		N018030 004
MILES	5GM/100ML;200MG/100ML		N018399 001
DEXTROSE 5% AND SODIUM CHLORIDE 0.225%	IN	PLASTIC CONTAINER	
ABBOTT	5GM/100ML;225MG/100ML		N019482 001 Oct 04, 1985

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DEXTROSE; SODIUM CHLORIDE

INJECTABLE; INJECTION

DEXTROSE 5% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER			
ABBOTT	5GM/100ML;300MG/100ML	N019486 001	Oct 04, 1985
MILES	5GM/100ML;300MG/100ML	N018501 001	
DEXTROSE 5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER			
B BRAUN	5GM/100ML;330MG/100ML	N018030 003	
DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER			
ABBOTT	5GM/100ML;450MG/100ML	N019484 001	Oct 04, 1985
B BRAUN	5GM/100ML;450MG/100ML	N018030 002	
MILES	5GM/100ML;450MG/100ML	N018400 001	
DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER			
ABBOTT	5GM/100ML;900MG/100ML	N019483 001	Oct 04, 1985
B BRAUN	5GM/100ML;900MG/100ML	N018026 001	
MILES	5GM/100ML;900MG/100ML	N018500 001	

DEXTROTHYROXINE SODIUM

TABLET; ORAL

CHOLOXIN			
ABBVIE	1MG	N012302 005	
	2MG	N012302 002	
	4MG	N012302 004	
	6MG	N012302 006	

DEZOCINE

INJECTABLE; INJECTION

DALGAN			
ASTRAZENECA	5MG/ML	N019082 001	Dec 29, 1989
	10MG/ML	N019082 002	Dec 29, 1989
	15MG/ML	N019082 003	Dec 29, 1989

DIATRIZOATE MEGLUMINE

INJECTABLE; INJECTION

ANGIOVIST 282			
BAYER HLTHCARE	60%	A087726 001	Sep 23, 1982
CARDIOGRAFIN			
BRACCO	85%	N011620 002	
DIATRIZOATE MEGLUMINE			
BRACCO	76%	N010040 017	
HYPAQUE			
GE HEALTHCARE	30%	N016403 002	
	60%	N016403 001	
RENO-60			
BRACCO	60%	N010040 016	
RENO-DIP			
BRACCO	30%	N010040 012	
UROVIST MEGLUMINE DIU/CT			
BAYER HLTHCARE	30%	A087739 001	Sep 23, 1982
SOLUTION; URETERAL			
RENO-30			
BRACCO	30%	N010040 021	
UROVIST CYSTO			
BAYER HLTHCARE	30%	A087729 001	Sep 23, 1982
UROVIST CYSTO PEDIATRIC			
BAYER HLTHCARE	30%	A087731 001	Sep 23, 1982
SOLUTION; URETHRAL			
HYPAQUE-CYSTO			
GE HEALTHCARE	30%	N016403 003	

DIATRIZOATE MEGLUMINE; DIATRIZOATE SODIUM

INJECTABLE; INJECTION

ANGIOVIST 292			
BAYER HLTHCARE	52%;8%	A087724 001	Sep 23, 1982
ANGIOVIST 370			
BAYER HLTHCARE	66%;10%	A087723 001	Sep 23, 1982
DIATRIZOATE-60			
INTL MEDICATION	52%;8%	A088166 001	Jun 17, 1983
HYPAQUE-76			
GE HEALTHCARE	66%;10%	A086505 001	
HYPAQUE-M, 75%			
GE HEALTHCARE	50%;25%	N010220 003	

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DIATRIZOATE MEGLUMINE; DIATRIZOATE SODIUM

INJECTABLE; INJECTION

HYPAQUE-M, 90%				
	GE HEALTHCARE	60%;30%	N010220	002
MD-60				
	MALLINCKRODT	52%;8%	A087074	001
MD-76				
	MALLINCKRODT	66%;10%	A087073	001
MD-76R				
+	LIEBEL-FLARSHEIM	66%;10%	N019292	001 Sep 29, 1989
RENOCAL-76				
	BRACCO	66%;10%	A089347	001 Jun 01, 1988
RENOGRAFIN-60				
	BRACCO	52%;8%	N010040	006
RENOGRAFIN-76				
+	BRACCO	66%;10%	N010040	001
RENOVIST				
	BRACCO	34.3%;35%	N010040	020
RENOVIST II				
	BRACCO	28.5%;29.1%	N010040	019
SOLUTION; ORAL, RECTAL				
GASTROVIST				
	BAYER HLTHCARE	66%;10%	A087728	001 Sep 23, 1982

DIATRIZOATE MEGLUMINE; IODIPAMIDE MEGLUMINE

SOLUTION; INTRAUTERINE

SINOGRAFIN				
+	BRACCO	52.7%;26.8%	N011324	002

DIATRIZOATE SODIUM

FOR SOLUTION; ORAL, RECTAL

HYPAQUE				
	GE HEALTHCARE	100%	N011386	001
INJECTABLE; INJECTION				
HYPAQUE				
	GE HEALTHCARE	25%	N009561	003
		50%	N009561	001
MD-50				
	MALLINCKRODT	50%	A087075	001
UROVIST SODIUM 300				
	BAYER HLTHCARE	50%	A087725	001 Sep 23, 1982
SOLUTION; ORAL, RECTAL				
HYPAQUE				
	GE HEALTHCARE	40%	N011386	003
SOLUTION; URETERAL				
HYPAQUE SODIUM 20%				
	GE HEALTHCARE	20%	N009561	002

DIAZEPAM

CAPSULE, EXTENDED RELEASE; ORAL

VALRELEASE				
	ROCHE	15MG	N018179	001
GEL; RECTAL				
DIASTAT				
+	BAUSCH	5MG/ML (5MG/ML) **	N020648	002 Jul 29, 1997
+		10MG/2ML (5MG/ML) **	N020648	003 Jul 29, 1997
+		15MG/3ML (5MG/ML) **	N020648	004 Jul 29, 1997
+		20MG/4ML (5MG/ML) **	N020648	005 Jul 29, 1997
INJECTABLE; INJECTION				
DIAZEPAM				
	ABRAXIS PHARM	5MG/ML	A070662	001 Jun 25, 1986
	HIKMA	5MG/ML (5MG/ML)	A070311	002 Dec 16, 1985
		5MG/ML	A070312	001 Dec 16, 1985
		5MG/ML	A071308	001 Jul 17, 1987
		5MG/ML	A071309	001 Jul 17, 1987
		5MG/ML	A071310	001 Jul 17, 1987
		10MG/2ML (5MG/ML)	A070311	003 Dec 16, 1985
	HOSPIRA	5MG/ML	A071584	001 Oct 13, 1987
	MARSAM PHARMS LLC	5MG/ML	A072371	001 Jan 29, 1993
	PARENTA PHARMS	5MG/ML	A076815	001 Apr 15, 2004
	US ARMY	5MG/ML **	N020124	001 Dec 05, 1990

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DIAZEPAM

INJECTABLE; INJECTION

DIAZEPAM

WARNER CHILCOTT	5MG/ML	A071613	001	Oct 22, 1987
	5MG/ML	A071614	001	Oct 22, 1987
WATSON LABS	5MG/ML	A070296	001	Feb 12, 1986
	5MG/ML	A070911	001	Aug 28, 1986
	5MG/ML	A070912	001	Aug 28, 1986
	5MG/ML	A070930	001	Dec 01, 1986
WATSON LABS INC	5MG/ML	A072370	001	Jan 29, 1993
	5MG/ML	A072397	001	Jan 29, 1993

DIZAC

PHARMACIA AND UPJOHN	5MG/ML **	N019287	001	Jun 18, 1993
----------------------	-----------	---------	-----	--------------

VALIUM

+ ROCHE	5MG/ML **	N016087	001	
---------	-----------	---------	-----	--

TABLET; ORAL

DIAZEPAM

ACTAVIS ELIZABETH	2MG	A070781	001	Mar 19, 1986
	5MG	A070706	001	Mar 19, 1986
	10MG	A070707	001	Mar 19, 1986
BARR	2MG	A070152	001	Nov 01, 1985
	10MG	A070154	001	Nov 01, 1985
CHARTWELL RX	2MG	A070302	001	Dec 20, 1985
	5MG	A070303	001	Dec 20, 1985
	10MG	A070304	001	Dec 20, 1985
DURAMED PHARMS BARR	2MG	A070894	001	Aug 27, 1986
	5MG	A070895	001	Aug 27, 1986
	10MG	A070896	001	Aug 27, 1986
ENDO OPERATIONS	2MG	A070228	002	Sep 26, 1985
	5MG	A070228	003	Sep 26, 1985
	10MG	A070228	001	Sep 26, 1985
FERNDALE LABS	2MG	A070903	001	Apr 01, 1987
	5MG	A070904	001	Apr 01, 1987
	10MG	A070905	001	Apr 01, 1987
HALSEY	2MG	A070987	001	Aug 15, 1986
	5MG	A070996	001	Aug 15, 1986
	10MG	A070956	001	Aug 15, 1986
IVAX SUB TEVA PHARMS	2MG	A070360	001	Sep 04, 1985
	5MG	A070361	001	Sep 04, 1985
	10MG	A070362	001	Sep 04, 1985
MARTEC USA LLC	10MG	A072402	001	Apr 25, 1989
PIONEER PHARMS	2MG	A070787	001	Aug 02, 1988
	5MG	A070788	001	Aug 02, 1988
	10MG	A070776	001	Aug 02, 1988
ROXANE	2MG	A070356	001	Jun 17, 1986
	5MG	A070357	001	Jun 17, 1986
	10MG	A070358	001	Jun 17, 1986
TEVA PHARMS	5MG	A070153	001	Nov 01, 1985
VIRTUS	2MG	A070462	001	Feb 25, 1986
	5MG	A070463	001	Feb 25, 1986
WARNER CHILCOTT	2MG	A070209	001	Sep 04, 1985
	5MG	A070210	001	Sep 04, 1985
	10MG	A070222	001	Sep 04, 1985
WATSON LABS	2MG	A070456	001	Nov 01, 1985
	5MG	A070457	001	Nov 01, 1985
	10MG	A070458	001	Nov 01, 1985

Q-PAM

QUANTUM PHARMICS	2MG	A070423	001	Dec 12, 1985
	2MG	A072431	001	Apr 29, 1988
	5MG	A070424	001	Dec 12, 1985
	5MG	A072432	001	Apr 29, 1988
	10MG	A070425	001	Dec 12, 1985
	10MG	A072433	001	Apr 29, 1988

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DIAZOXIDECAPSULE; ORAL
PROGLYCEMTEVA BRANDED PHARM 50MG N017425 001
100MG N017425 002INJECTABLE; INJECTION
DIAZOXIDEABRAXIS PHARM 15MG/ML A071519 001 Aug 26, 1987
HYPERSTAT
SCHERING 15MG/ML N016996 001DIBUCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

HEAVY SOLUTION NUPERCALINE
NOVARTIS 2.5MG/ML N006203 001DICHLORPHENAMIDE

TABLET; ORAL

DARANIDE
+ XERIS 50MG ** N011366 001DICLOFENAC

CAPSULE; ORAL

ZORVOLEX
+ ZYLA 18MG ** N204592 001 Oct 18, 2013
+ 35MG ** N204592 002 Oct 18, 2013DICLOFENAC POTASSIUM

TABLET; ORAL

CATAFLAM
+ NOVARTIS 25MG ** N020142 001 Nov 24, 1993
+ 50MG ** N020142 002 Nov 24, 1993
DICLOFENAC POTASSIUM
CHARTWELL RX 50MG A075582 001 Feb 23, 2001
SUN PHARM INDUSTRIES 50MG A075470 001 Feb 21, 2002
WATSON LABS TEVA 50MG A075152 001 Nov 27, 1998DICLOFENAC SODIUM

GEL; TOPICAL

DICLOFENAC SODIUM
HIKMA 1% A209484 001 Nov 21, 2018
SOLARAZE
+ FOUGERA PHARMS 3% ** N021005 001 Oct 16, 2000

SOLUTION; INTRAVENOUS

DICLOFENAC SODIUM
RISING 37.5MG/ML (37.5MG/ML) A208786 001 Jun 18, 2019
DYLOJECT
+ JAVELIN PHARMS INC 37.5MG/ML (37.5MG/ML) ** N022396 001 Dec 23, 2014

SOLUTION; TOPICAL

DICLOFENAC SODIUM
APOTEX 1.5% A202027 001 May 27, 2014
PAI HOLDINGS PHARM 1.5% A202769 001 Jul 08, 2015
RISING 1.5% A206715 001 Aug 07, 2017
TWI PHARMS 1.5% A202393 001 Nov 24, 2014
PENNSAID
+ HORIZON 2% ** N204623 001 Jan 16, 2014
+ NUVO PHARMS INC 1.5% ** N020947 001 Nov 04, 2009

SOLUTION/DROPS; OPHTHALMIC

DICLOFENAC SODIUM
FALCON PHARMS 0.1% N020809 001 May 04, 1998
SCIEGEN PHARMS INC 0.1% A077845 001 Apr 17, 2008
VOLTAREN
+ NOVARTIS 0.1% N020037 001 Mar 28, 1991

TABLET, DELAYED RELEASE; ORAL

DICLOFENAC SODIUM
AUROBINDO PHARMA USA 50MG A075281 002 Feb 12, 2002
75MG A075281 003 Feb 12, 2002
CHARTWELL RX 25MG A074376 001 Sep 28, 1995
50MG A074376 002 Sep 28, 1995
75MG A074394 001 Nov 30, 1995
MICRO LABS 50MG A074986 001 Feb 26, 1999
75MG A074986 002 Feb 26, 1999

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DICLOFENAC SODIUM

TABLET, DELAYED RELEASE;ORAL

DICLOFENAC SODIUM

PLIVA	50MG	A074432 002	Jul 29, 1999
	75MG	A074432 003	Jul 29, 1999
ROXANE	25MG	A074391 001	Jun 29, 1995
	50MG	A074391 002	Jun 29, 1995
	75MG	A074391 003	Jun 29, 1995
TEVA	50MG	A074723 001	Mar 30, 1999
	75MG	A074390 001	Aug 15, 1996
TEVA PHARMS	25MG	A074459 001	Jun 25, 1997
	50MG	A074459 002	Jun 25, 1997
	75MG	A074459 003	Jun 25, 1997
VOLTAREN			
+ NOVARTIS	25MG **	N019201 001	Jul 28, 1988
+	50MG **	N019201 002	Jul 28, 1988
+	75MG **	N019201 003	Jul 28, 1988

TABLET, EXTENDED RELEASE;ORAL

DICLOFENAC SODIUM

ACTAVIS ELIZABETH	100MG	A075910 001	Jan 07, 2002
AUROBINDO PHARMA USA	100MG	A076152 001	Dec 13, 2001
VOLTAREN-XR			
+ NOVARTIS	100MG **	N020254 001	Mar 08, 1996

DICLOFENAC SODIUM; MISOPROSTOL

TABLET, DELAYED RELEASE;ORAL

DICLOFENAC SODIUM AND MISOPROSTOL

EXELA HOLDINGS	50MG;0.2MG	A200540 001	Mar 14, 2014
	75MG;0.2MG	A200540 002	Mar 14, 2014
SANDOZ	50MG;0.2MG	A200158 001	May 09, 2013
	75MG;0.2MG	A200158 002	May 09, 2013
ZYDUS PHARMS	50MG;0.2MG	A206771 001	Jun 12, 2023
	75MG;0.2MG	A206771 002	Jun 12, 2023

DICLOXACILLIN SODIUM

CAPSULE;ORAL

DICLOXACILLIN SODIUM

CHARTWELL RX	EQ 125MG BASE	A061454 002	
	EQ 250MG BASE	A061454 001	
	EQ 500MG BASE	A061454 003	
DYCILL			
GLAXOSMITHKLINE	EQ 250MG BASE	A060254 002	
	EQ 250MG BASE	A062238 001	
	EQ 500MG BASE	A060254 003	
	EQ 500MG BASE	A062238 002	
PATHOCIL			
+ WYETH AYERST	EQ 250MG BASE **	N050011 002	
+	EQ 500MG BASE **	N050011 003	Mar 28, 1983

FOR SUSPENSION;ORAL

DICLOXACILLIN SODIUM

APOTHECON	EQ 62.5MG BASE/5ML	A061455 001	
DYNAPEN			
APOTHECON	EQ 62.5MG BASE/5ML	N050337 002	
PATHOCIL			
WYETH AYERST	EQ 62.5MG BASE/5ML	N050092 001	

DICUMAROL

CAPSULE;ORAL

DICUMAROL

LILLY	25MG	N005509 003	
	50MG	N005509 001	

TABLET;ORAL

DICUMAROL

ABBVIE	25MG	N005545 003	
	50MG	N005545 004	
	100MG	N005545 005	

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DICYCLOMINE HYDROCHLORIDE

CAPSULE; ORAL

BENTYL

+	ALLERGAN	10MG **	N007409 003	Oct 15, 1984
---	----------	---------	-------------	--------------

DICYCLOMINE HYDROCHLORIDE

	PIONEER PHARMS	10MG	A089361 001	Jan 10, 1989
--	----------------	------	-------------	--------------

	SUN PHARM INDUSTRIES	10MG	A084505 001	Oct 21, 1986
--	----------------------	------	-------------	--------------

	WATSON LABS	10MG	A083179 001	Feb 12, 1986
--	-------------	------	-------------	--------------

INJECTABLE; INJECTION

BENTYL

+	ABBVIE	10MG/ML	N008370 001	Oct 15, 1984
---	--------	---------	-------------	--------------

BENTYL PRESERVATIVE FREE

+	ABBVIE	10MG/ML **	N008370 002	Oct 15, 1984
---	--------	------------	-------------	--------------

DICYCLOMINE HYDROCHLORIDE

	DR REDDYS	10MG/ML	A080614 001	Feb 11, 1986
--	-----------	---------	-------------	--------------

	HIKMA	10MG/ML	A210788 001	Feb 11, 2019
--	-------	---------	-------------	--------------

	PRAXGEN PHARMS	10MG/ML	A212058 001	Apr 26, 2019
--	----------------	---------	-------------	--------------

	RENEW PHARMS	10MG/ML	A207084 001	May 04, 2018
--	--------------	---------	-------------	--------------

SYRUP; ORAL

BENTYL

+	APTALIS PHARMA US	10MG/5ML **	N007961 002	Oct 15, 1984
---	-------------------	-------------	-------------	--------------

DICYCLOMINE HYDROCHLORIDE

	ALPHARMA US PHARMS	10MG/5ML	A084479 001	
--	--------------------	----------	-------------	--

TABLET; ORAL

BENTYL

+	ALLERGAN	20MG **	N007409 001	Oct 15, 1984
---	----------	---------	-------------	--------------

DICYCLOMINE HYDROCHLORIDE

	PIONEER PHARMS	20MG	A088585 001	Aug 20, 1986
--	----------------	------	-------------	--------------

	SUN PHARM INDUSTRIES	20MG	A084600 001	Jul 29, 1985
--	----------------------	------	-------------	--------------

	WATSON LABS	20MG	A084361 001	Feb 06, 1986
--	-------------	------	-------------	--------------

DIDANOSINE

CAPSULE, DELAYED REL PELLETS; ORAL

DIDANOSINE

	AUROBINDO PHARMA	125MG	A090094 001	Sep 24, 2008
--	------------------	-------	-------------	--------------

		200MG	A090094 002	Sep 24, 2008
--	--	-------	-------------	--------------

		250MG	A090094 003	Sep 24, 2008
--	--	-------	-------------	--------------

		400MG	A090094 004	Sep 24, 2008
--	--	-------	-------------	--------------

	BARR	200MG	A077167 001	Dec 03, 2004
--	------	-------	-------------	--------------

		250MG	A077167 002	Dec 03, 2004
--	--	-------	-------------	--------------

		400MG	A077167 003	Dec 03, 2004
--	--	-------	-------------	--------------

	NORVIUM BIOSCIENCE	125MG	A090788 001	Apr 08, 2010
--	--------------------	-------	-------------	--------------

		200MG	A090788 002	Apr 08, 2010
--	--	-------	-------------	--------------

		250MG	A090788 003	Apr 08, 2010
--	--	-------	-------------	--------------

		400MG	A090788 004	Apr 08, 2010
--	--	-------	-------------	--------------

VIDEX EC

+	BRISTOL MYERS SQUIBB	125MG **	N021183 001	Oct 31, 2000
---	----------------------	----------	-------------	--------------

+		200MG **	N021183 002	Oct 31, 2000
---	--	----------	-------------	--------------

+		250MG **	N021183 003	Oct 31, 2000
---	--	----------	-------------	--------------

+		400MG **	N021183 004	Oct 31, 2000
---	--	----------	-------------	--------------

FOR SOLUTION; ORAL

DIDANOSINE

	AUROBINDO PHARMA	10MG/ML	A078112 001	Mar 08, 2007
--	------------------	---------	-------------	--------------

VIDEX

+	BRISTOL	10MG/ML	N020156 001	Oct 09, 1991
---	---------	---------	-------------	--------------

	BRISTOL MYERS SQUIBB	100MG/PACKET	N020155 003	Oct 09, 1991
--	----------------------	--------------	-------------	--------------

		167MG/PACKET	N020155 004	Oct 09, 1991
--	--	--------------	-------------	--------------

		250MG/PACKET	N020155 005	Oct 09, 1991
--	--	--------------	-------------	--------------

		375MG/PACKET	N020155 006	Oct 09, 1991
--	--	--------------	-------------	--------------

TABLET, CHEWABLE; ORAL

VIDEX

+	BRISTOL MYERS SQUIBB	25MG **	N020154 002	Oct 09, 1991
---	----------------------	---------	-------------	--------------

+		50MG **	N020154 003	Oct 09, 1991
---	--	---------	-------------	--------------

+		100MG **	N020154 004	Oct 09, 1991
---	--	----------	-------------	--------------

+		150MG **	N020154 005	Oct 09, 1991
---	--	----------	-------------	--------------

+		200MG **	N020154 006	Oct 28, 1999
---	--	----------	-------------	--------------

TABLET, FOR SUSPENSION; ORAL

DIDANOSINE

	AUROBINDO	100MG	A077275 001	Aug 14, 2012
--	-----------	-------	-------------	--------------

		150MG	A077275 002	Aug 14, 2012
--	--	-------	-------------	--------------

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DIDANOSINETABLET, FOR SUSPENSION;ORAL
DIDANOSINE

200MG

A077275 003 Aug 14, 2012

DIENESTROLCREAM;VAGINAL
DIENESTROL

ORTHO MCNEIL PHARM 0.01%

N006110 005

DV

SANOFI AVENTIS US 0.01%

A083518 001

ESTRAGUARD

SOLVAY 0.01%

A084436 001

SUPPOSITORY;VAGINAL

DV

SANOFI AVENTIS US 0.7MG

A083517 001

DIETHYLCARBAMAZINE CITRATETABLET;ORAL
HETRAZAN

LEDERLE 50MG

N006459 001

DIETHYLPROPION HYDROCHLORIDE

TABLET;ORAL

DIETHYLPROPION HYDROCHLORIDE

CHARTWELL RX 25MG

A088267 001 Aug 25, 1983

25MG

A088268 001 Aug 25, 1983

EPIC PHARMA LLC 25MG

A040828 001 Nov 05, 2008

SANDOZ 25MG

A085916 001

TEVA 25MG

A088642 001 Sep 20, 1984

UCB INC 25MG

A085544 001

WATSON LABS 25MG

A085741 001

TENUATE

+ NOSTRUM LABS INC 25MG **

N011722 002

SANOFI AVENTIS US 25MG

N017668 001

TEPANIL

3M 25MG

N011673 001

TABLET, EXTENDED RELEASE;ORAL

TENUATE

SANOFI AVENTIS US 75MG

N017669 001

TENUATE DOSPAN

+ NOSTRUM LABS INC 75MG **

N012546 001

TEPANIL TEN-TAB

3M 75MG

N017956 001

DIETHYLSTILBESTROLINJECTABLE;INJECTION
STILBESTROL

BRISTOL MYERS SQUIBB 0.2MG/ML

N004056 003

0.5MG/ML

N004056 004

1MG/ML

N004056 005

5MG/ML

N004056 006

SUPPOSITORY;VAGINAL

DIETHYLSTILBESTROL

LILLY 0.1MG

N004040 001

0.5MG

N004040 002

STILBESTROL

BRISTOL MYERS SQUIBB 0.1MG

N004056 001

0.5MG

N004056 002

TABLET;ORAL

DIETHYLSTILBESTROL

LILLY 0.1MG

N004041 002

0.5MG

N004041 003

1MG

N004041 004

5MG

N004041 005

STILBESTROL

TABLICAPS 0.5MG

A083004 001

1MG

A083002 001

5MG

A083006 001

STILBETIN

BRISTOL MYERS SQUIBB 0.1MG

N004056 007

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DIETHYLSTILBESTROLTABLET; ORAL
STILBETIN

0.25MG	N004056 017
0.5MG	N004056 008
1MG	N004056 009
5MG	N004056 010

TABLET, DELAYED RELEASE; ORAL
DIETHYLSTILBESTROL

LILLY	0.1MG	N004039 002
	0.25MG	N004039 005
	0.5MG	N004039 003
	1MG	N004039 004
	5MG	N004039 006

STILBESTROL

TABLICAPS	0.5MG	A083003 001
	1MG	A083005 001
	5MG	A083007 001

STILBETIN

BRISTOL MYERS SQUIBB	0.1MG	N004056 011
	0.5MG	N004056 012
	1MG	N004056 013
	5MG	N004056 014

DIETHYLSTILBESTROL DIPHOSPHATE

INJECTABLE; INJECTION

STILPHOSTROL

BAYER PHARMS	250MG/5ML	N010010 001
--------------	-----------	-------------

TABLET; ORAL

STILPHOSTROL

BAYER PHARMS	50MG	N010010 002
--------------	------	-------------

DIFLORASONE DIACETATE

CREAM; TOPICAL

DIFLORASONE DIACETATE

ANI PHARMS	0.05%	A076263 001	Dec 20, 2002
AVONDALE PHARMS	0.05%	A075187 001	Mar 30, 1998

FLORONE

PFIZER	0.05% **	N017741 001
--------	----------	-------------

FLORONE E

PFIZER	0.05%	N019259 001	Aug 28, 1985
--------	-------	-------------	--------------

PSORCON

+ TARO PHARMS NORTH	0.05% **	N020205 001	Nov 20, 1992
---------------------	----------	-------------	--------------

OINTMENT; TOPICAL

DIFLORASONE DIACETATE

SCIEGEN PHARMS INC	0.05%	A206572 001	Jul 24, 2015
--------------------	-------	-------------	--------------

PSORCON

+ PFIZER	0.05%	N019260 001	Aug 28, 1985
----------	-------	-------------	--------------

PSORCON E

PFIZER	0.05%	N017994 001
--------	-------	-------------

DIFLUNISAL

TABLET; ORAL

DIFLUNISAL

ANI PHARMS	500MG	A074604 001	Jun 10, 1996
------------	-------	-------------	--------------

DASTECH INTL	250MG	A073562 001	Nov 27, 1992
--------------	-------	-------------	--------------

	500MG	A073563 001	Nov 27, 1992
--	-------	-------------	--------------

PUREPAC PHARM	250MG	A074285 001	May 07, 1996
---------------	-------	-------------	--------------

	500MG	A074285 002	May 07, 1996
--	-------	-------------	--------------

TEVA	250MG	A073679 001	Jul 31, 1992
------	-------	-------------	--------------

WATSON LABS	250MG	A074400 001	Jul 17, 1997
-------------	-------	-------------	--------------

	500MG	A074400 002	Jul 17, 1997
--	-------	-------------	--------------

DOLOBID

+ MERCK	250MG **	N018445 001	Apr 19, 1982
---------	----------	-------------	--------------

+	500MG **	N018445 002	Apr 19, 1982
---	----------	-------------	--------------

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DIGITOXIN

INJECTABLE; INJECTION

CRYSTODIGIN

LILLY

0.2MG/ML

A084100 005

DIGOXIN

CAPSULE; ORAL

LANOXICAPS

GLAXOSMITHKLINE LLC

0.05MG

N018118 002 Jul 26, 1982

0.1MG

N018118 003 Jul 26, 1982

0.15MG

N018118 004 Sep 24, 1984

0.2MG

N018118 001 Jul 26, 1982

INJECTABLE; INJECTION

DIGOXIN

ABRAXIS PHARM

0.25MG/ML

A083217 001

HOSPIRA

0.25MG/ML

A040093 001 May 16, 1996

0.25MG/ML

A040206 001 Aug 28, 1998

WYETH AYERST

0.25MG/ML

A084386 001

DIGOXIN PEDIATRIC

HOSPIRA

0.1MG/ML

A040092 001 Apr 25, 1996

TABLET; ORAL

DIGOXIN

IMPAX LABS

0.125MG

A078556 001 Jul 20, 2009

0.25MG

A078556 002 Jul 20, 2009

LANOXIN

+ ADVANZ PHARMA

0.1875MG

N020405 003 Sep 30, 1997

0.375MG

N020405 005 Sep 30, 1997

0.5MG

N020405 006 Sep 30, 1997

DIHYDROERGOTAMINE MESYLATE

INJECTABLE; INJECTION

D.H.E. 45

+ BAUSCH

1MG/ML **

N005929 001

SPRAY, METERED; NASAL

DIHYDROERGOTAMINE MESYLATE

AMNEAL

0.5MG/SPRAY

A214105 001 Jan 04, 2022

DIHYDROERGOTAMINE MESYLATE; HEPARIN SODIUM; LIDOCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

EMBOLEX

NOVARTIS

0.5MG/0.5ML; 2,500

N018885 001 Nov 30, 1984

UNITS/0.5ML; 5.33MG/0.5ML

0.5MG/0.7ML; 5,000

N018885 002 Nov 30, 1984

UNITS/0.7ML; 7.46MG/0.7ML

DILTIAZEM HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

CARDIZEM SR

+ BIOVAIL

60MG **

N019471 001 Jan 23, 1989

+

90MG **

N019471 002 Jan 23, 1989

+

120MG **

N019471 003 Jan 23, 1989

+

180MG **

N019471 004 Jan 23, 1989

DILACOR XR

+ ALLERGAN

120MG **

N020092 001 May 29, 1992

+

180MG **

N020092 002 May 29, 1992

+

240MG **

N020092 003 May 29, 1992

DILT-CD

APOTEX

120MG

A076151 001 May 20, 2004

180MG

A076151 002 May 20, 2004

240MG

A076151 003 May 20, 2004

300MG

A076151 004 May 20, 2004

DILTIAZEM HYDROCHLORIDE

ACTAVIS LABS FL INC

120MG

A074852 001 Oct 10, 1997

180MG

A074852 002 Oct 10, 1997

240MG

A074852 003 Oct 10, 1997

BIOVAIL

60MG

A074845 001 Sep 15, 1999

90MG

A074845 002 Sep 15, 1999

120MG

A074845 003 Sep 15, 1999

120MG

N020939 001 Jan 28, 2000

180MG

N020939 002 Jan 28, 2000

240MG

N020939 003 Jan 28, 2000

300MG

N020939 004 Jan 28, 2000

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DILTIAZEM HYDROCHLORIDECAPSULE, EXTENDED RELEASE;ORAL
DILTIAZEM HYDROCHLORIDE

	360MG	N020939 005	Sep 14, 2001
	420MG	N020939 006	Sep 14, 2001
ENDO OPERATIONS	360MG	A209766 001	May 30, 2018
NESHER PHARMS	120MG	A076563 002	Sep 12, 2006
	180MG	A076563 003	Sep 12, 2006
	240MG	A076563 004	Sep 12, 2006
	300MG	A076563 005	Sep 12, 2006
	360MG	A076563 006	Sep 12, 2006
	420MG	A076563 001	Sep 12, 2006
NORVIUM BIOSCIENCE	120MG	A075124 002	Mar 18, 1998
	180MG	A075124 003	Mar 18, 1998
	240MG	A075124 001	Mar 18, 1998
SUN PHARM	120MG	A090421 001	Nov 15, 2010
	120MG	A090492 001	Oct 28, 2011
	180MG	A090421 002	Nov 15, 2010
	180MG	A090492 002	Oct 28, 2011
	240MG	A090421 003	Nov 15, 2010
	240MG	A090492 003	Oct 28, 2011
	300MG	A090421 004	Nov 15, 2010
	300MG	A090492 004	Oct 28, 2011
	360MG	A090421 005	Nov 15, 2010
	360MG	A090492 005	Oct 28, 2011
TEVA	60MG	A074079 001	Nov 30, 1993
	90MG	A074079 002	Nov 30, 1993
	120MG	A074079 003	Nov 30, 1993
DILTIZAC			
APOTEX	120MG	A076395 001	Feb 01, 2006
	180MG	A076395 002	Feb 01, 2006
	240MG	A076395 003	Feb 01, 2006
	300MG	A076395 004	Feb 01, 2006
	360MG	A076395 005	Feb 01, 2006
TAZTIA XT			
ACTAVIS LABS FL INC	120MG	A075401 001	Apr 10, 2003
	180MG	A075401 002	Apr 10, 2003
	240MG	A075401 003	Apr 10, 2003
	300MG	A075401 004	Apr 10, 2003
	360MG	A075401 005	Apr 10, 2003
INJECTABLE; INJECTION			
CARDIZEM			
BIOVAIL	100MG/VIAL **	N020792 001	Sep 05, 1997
+ BIOVAIL LABS INTL	5MG/ML **	N020027 001	Oct 24, 1991
+	25MG/VIAL **	N020027 003	Aug 18, 1995
DILTIAZEM HYDROCHLORIDE			
DR REDDYS	5MG/ML	A074894 001	Aug 26, 1997
HIKMA FARMACEUTICA	5MG/ML	A202651 001	Aug 09, 2012
HOSPIRA	5MG/ML	A075004 001	Feb 16, 2000
	5MG/ML	A075106 001	Apr 29, 1999
INTL MEDICATION	5MG/ML	A075749 001	Nov 21, 2001
MYLAN LABS LTD	5MG/ML	A075375 001	Sep 30, 1999
SOLUTION; INTRAVENOUS			
DILTIAZEM HYDROCHLORIDE IN DEXTROSE 5%			
+ EXELA PHARMA	125MG/125ML (1MG/ML) **	N215252 001	Oct 28, 2021
+	250MG/250ML (1MG/ML) **	N215252 002	Oct 28, 2021
TABLET; ORAL			
DILTIAZEM HYDROCHLORIDE			
APOTHECON	30MG	A074051 001	Mar 31, 1993
	60MG	A074051 002	Mar 31, 1993
	90MG	A074051 003	Mar 31, 1993
	120MG	A074051 004	Mar 31, 1993
CHARTWELL MOLECULES	30MG	A074093 001	Nov 05, 1992
	60MG	A074093 002	Nov 05, 1992
	90MG	A074093 003	Nov 05, 1992
	120MG	A074093 004	Nov 05, 1992
EDENBRIDGE PHARMS	30MG	A211596 001	Nov 18, 2019
	60MG	A211596 002	Nov 18, 2019
	90MG	A211596 003	Nov 18, 2019

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DILTIAZEM HYDROCHLORIDE

TABLET; ORAL

DILTIAZEM HYDROCHLORIDE

	120MG	A211596 004	Nov 18, 2019
IVAX SUB TEVA PHARMS	30MG	A074168 001	Mar 03, 1995
	60MG	A074168 002	Mar 03, 1995
	90MG	A074168 003	Mar 03, 1995
	120MG	A074168 004	Mar 03, 1995
RISING	30MG	A072838 004	Nov 05, 1992
	60MG	A072838 003	Nov 05, 1992
	90MG	A072838 002	Nov 05, 1992
	120MG	A072838 001	Nov 05, 1992
TEVA	30MG	A074084 001	Feb 25, 1994
	60MG	A074084 002	Feb 25, 1994
TEVA PHARMS	30MG	A074067 001	Nov 05, 1992
	60MG	A074067 002	Nov 05, 1992
	90MG	A074067 003	Nov 05, 1992
	120MG	A074067 004	Nov 05, 1992

DILTIAZEM MALATE

TABLET, EXTENDED RELEASE; ORAL

TIAMATE

MERCK	EQ 120MG HYDROCHLORIDE	N020506 001	Oct 04, 1996
	EQ 180MG HYDROCHLORIDE	N020506 002	Oct 04, 1996
	EQ 240MG HYDROCHLORIDE	N020506 003	Oct 04, 1996

DILTIAZEM MALATE; ENALAPRIL MALEATE

TABLET, EXTENDED RELEASE; ORAL

TECZEM

BIOVAIL	EQ 180MG HYDROCHLORIDE; 5MG	N020507 001	Oct 04, 1996
---------	-----------------------------	-------------	--------------

DIMENHYDRINATE

INJECTABLE; INJECTION

DIMENHYDRINATE

BAXTER HLTHCARE	50MG/ML	A084767 001	
WATSON LABS	50MG/ML	A083531 001	
+ WATSON LABS TEVA	50MG/ML	A080615 001	
WYETH AYERST	50MG/ML	A084316 001	

LIQUID; ORAL

DIMENHYDRINATE

ALRA	12.5MG/4ML	A080715 001	
------	------------	-------------	--

TABLET; ORAL

DIMENHYDRINATE

HEATHER	50MG	A080841 001	
NEXGEN PHARMA INC	50MG	A085985 001	
+ WATSON LABS	50MG	A085166 001	

DIMETHYL FUMARATE

CAPSULE, DELAYED RELEASE; ORAL

DIMETHYL FUMARATE

SAWAI USA	120MG	A210285 001	Dec 21, 2021
	240MG	A210285 002	Dec 21, 2021
ZYDUS PHARMS	120MG	A210538 001	Sep 24, 2020
	240MG	A210538 002	Sep 24, 2020

DIMETHYL SULFOXIDE

SOLUTION; INTRAVESICAL

DIMETHYL SULFOXIDE

MYLAN INSTITUTIONAL	50%	A076185 001	Nov 29, 2002
---------------------	-----	-------------	--------------

DIMYRISTOYL LECITHIN; PERFLEXANE

INJECTABLE; INTRAVENOUS

IMAGENT

VESSELON SPV LLC	0.92MG/VIAL; 0.092MG/VIAL	N021191 001	May 31, 2002
------------------	---------------------------	-------------	--------------

DINOPROST TROMETHAMINE

INJECTABLE; INJECTION

PROSTIN F2 ALPHA

PHARMACIA AND UPJOHN	EQ 5MG BASE/ML	N017434 001	
----------------------	----------------	-------------	--

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DINOPROSTONE

SUPPOSITORY; VAGINAL
PROSTIN E2

+ PFIZER 20MG N017810 001

DIPHEMANIL METHYLSULFATE

TABLET; ORAL
PRANTAL

SCHERING 100MG N008114 004

DIPHENHYDRAMINE HYDROCHLORIDE

CAPSULE; ORAL

BENADRYL

MCNEIL CONS 25MG N005845 007

50MG N005845 001

DIPHENHYDRAMINE HYDROCHLORIDE

ALRA 25MG A080519 004

50MG A080519 003

ANABOLIC 50MG A083275 001

ELKINS SINN 25MG A085701 001

50MG A085701 002

FOSUN PHARMA 25MG A080832 001

25MG A080845 002

50MG A080832 002

50MG A080845 001

HALSEY 50MG A087914 001 Jun 04, 1984

HEATHER 25MG A084524 001

50MG A083953 001

HERITAGE PHARMA 50MG A080727 001

50MG A080738 001

HIKMA INTL PHARMS 50MG A083567 001

IMPAX LABS 25MG A080807 001

50MG A080807 002

IVAX SUB TEVA PHARMS 25MG A080762 001

50MG A080762 002

LANNETT 25MG A080868 001

50MG A080868 002

LEDERLE 25MG A086874 001

50MG A086875 001

LNK 25MG A087977 001 Jan 27, 1983

50MG A087978 001 Jan 27, 1983

MK LABS 25MG A083087 001

50MG A083087 002

MUTUAL PHARM 25MG A084506 001

NEWTRON PHARMS 25MG A086543 001

50MG A086544 001

NEXGEN PHARMA INC 25MG A083634 001

PERRIGO 25MG A083061 001

50MG A083061 002

PIONEER PHARMS 25MG A089101 001 Dec 20, 1985

50MG A088880 001 Dec 20, 1985

PUREPAC PHARM 25MG A085156 001

50MG A085150 001

PVT FORM 25MG A083027 001

50MG A083027 002

ROXANE 50MG A080635 001

SUN PHARM INDUSTRIES 25MG A089488 001 Jan 02, 1987

50MG A089489 001 Jan 02, 1987

SUPERPHARM 25MG A089040 001 May 15, 1985

50MG A089041 001 May 15, 1985

TEVA 25MG A085874 001

50MG A085874 002

VALEANT PHARM INTL 25MG A080596 001

50MG A080592 001

VANGARD 25MG A088034 001 Oct 27, 1982

50MG A087630 001

WATSON LABS 25MG A080728 001

25MG A083797 001

25MG A085138 001

50MG A083797 002

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DIPHENHYDRAMINE HYDROCHLORIDE

CAPSULE; ORAL

DIPHENHYDRAMINE HYDROCHLORIDE

	50MG	A085083	001	
WHITEWORTH TOWN PLSN	25MG	A083441	001	
	50MG	A080800	001	

ELIXIR; ORAL

BELIX

HALSEY	12.5MG/5ML	A086586	001	Oct 03, 1983
--------	------------	---------	-----	--------------

BENADRYL

MCNEIL CONS	12.5MG/5ML	N005845	004	
-------------	------------	---------	-----	--

DIBENIL

CENCI	12.5MG/5ML	A088304	001	Dec 16, 1983
-------	------------	---------	-----	--------------

DIPHEN

USL PHARMA	12.5MG/5ML	A084640	001	
------------	------------	---------	-----	--

DIPHENHYDRAMINE HYDROCHLORIDE

BUNDY	12.5MG/5ML	A083674	001	
CENCI	12.5MG/5ML	A087941	001	Dec 17, 1982
KV PHARM	12.5MG/5ML	A085621	001	
LANNETT	12.5MG/5ML	A080939	002	
LEDERLE	12.5MG/5ML	A086937	001	
MK LABS	12.5MG/5ML	A083088	002	
NASKA	12.5MG/5ML	A088680	001	May 31, 1985
PERRIGO	12.5MG/5ML	A083063	001	
PUREPAC PHARM	12.5MG/5ML	A083237	001	Jan 25, 1982
PVT FORM	12.5MG/5ML	A085287	001	
ROXANE	12.5MG/5ML	A080643	001	

HYDRAMINE

ALPHARMA US PHARMS	12.5MG/5ML	A080763	002	
--------------------	------------	---------	-----	--

INJECTABLE; INJECTION

BENADRYL

MCNEIL CONS	10MG/ML	N006146	001	
+	50MG/ML **	N006146	002	

BENADRYL PRESERVATIVE FREE

+	MCNEIL CONS	50MG/ML **	N009486	001
---	-------------	------------	---------	-----

DIPHENHYDRAMINE HYDROCHLORIDE

BEL MAR	10MG/ML	A080822	001	
DR REDDYS	10MG/ML	A080873	001	
	50MG/ML	A080873	002	
HIKMA	50MG/ML	A083183	001	
HOSPIRA	50MG/ML	A040140	001	Nov 20, 1998
LYPHOMED	10MG/ML	A087066	001	
WATSON LABS	10MG/ML	A083533	001	
WYETH AYERST	50MG/ML	A080577	001	

DIPHENHYDRAMINE HYDROCHLORIDE PRESERVATIVE FREE

ABRAXIS PHARM	50MG/ML	A080586	002	
DR REDDYS	50MG/ML	A080873	003	
INTL MEDICATION	50MG/ML	A084094	001	

SYRUP; ORAL

ANTITUSSIVE

PERRIGO	12.5MG/5ML	A071292	001	Apr 10, 1987
---------	------------	---------	-----	--------------

BELDIN

HALSEY	12.5MG/5ML	A089179	001	Jun 05, 1986
--------	------------	---------	-----	--------------

BENYLIN

PARKE DAVIS	12.5MG/5ML	N006514	004	
-------------	------------	---------	-----	--

DIPHEN

MORTON GROVE	12.5MG/5ML	A070118	001	Oct 01, 1985
--------------	------------	---------	-----	--------------

DIPHENHYDRAMINE HYDROCHLORIDE

ALPHARMA US PHARMS	12.5MG/5ML	A070497	001	Apr 25, 1989
CUMBERLAND SWAN	12.5MG/5ML	A073611	001	Aug 20, 1992
SCIEGEN PHARMS INC	12.5MG/5ML	A072416	001	Sep 28, 1990

HYDRAMINE

ALPHARMA US PHARMS	12.5MG/5ML	A070205	001	Jan 28, 1986
--------------------	------------	---------	-----	--------------

SILPHEN

LANNETT CO INC	12.5MG/5ML	A072646	001	Feb 27, 1992
----------------	------------	---------	-----	--------------

VICKS FORMULA 44

WARNER CHILCOTT	12.5MG/5ML	A070524	001	Jan 14, 1987
-----------------	------------	---------	-----	--------------

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DIPHENHYDRAMINE HYDROCHLORIDE; NAPROXEN SODIUM

TABLET; ORAL

NAPROXEN SODIUM AND DIPHENHYDRAMINE HYDROCHLORIDE

P AND L

25MG; 220MG

A207597 001 Jan 25, 2019

DIPHENHYDRAMINE HYDROCHLORIDE; PSEUDOEPHEDRINE HYDROCHLORIDE

SOLUTION; ORAL

BENYLIN

PARKE DAVIS

12.5MG/5ML; 30MG/5ML

N019014 001 Jun 11, 1985

DIPHENIDOL HYDROCHLORIDE

TABLET; ORAL

VONTROL

GLAXOSMITHKLINE

EQ 25MG BASE

N016033 001

DIPHENYLPYRALINE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

HISPRIL

GLAXOSMITHKLINE

5MG

N011945 001

DIPIVEFRIN HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

AKPRO

EPIC PHARMA LLC

0.1%

A074382 001 Sep 29, 1995

DIPIVEFRIN HYDROCHLORIDE

BAUSCH AND LOMB

0.1%

A074188 001 May 19, 1995

FALCON PHARMS

0.1%

A073636 001 Jun 30, 1994

PROPINE

ALLERGAN

0.1%

N018239 001

DIPYRIDAMOLE

INJECTABLE; INJECTION

DIPYRIDAMOLE

DR REDDYS

5MG/ML

A074952 001 Nov 26, 1997

EUGIA PHARMA SPECLTS

5MG/ML

A075769 001 Nov 27, 2002

FRESENIUS KABI USA

5MG/ML

A074956 001 Sep 30, 1998

HOSPIRA

5MG/ML

A074601 001 Dec 19, 1997

IV PERSANTINE

+ BOEHRINGER INGELHEIM

5MG/ML **

N019817 001 Dec 13, 1990

TABLET; ORAL

DIPYRIDAMOLE

ANI PHARMS

25MG

A086944 002 Apr 16, 1991

50MG

A086944 001 Feb 25, 1992

75MG

A086944 003 Feb 25, 1992

CHARTWELL RX

25MG

A040898 001 Apr 23, 2008

50MG

A040898 002 Apr 23, 2008

75MG

A040898 003 Apr 23, 2008

GLENMARK GENERICS

25MG

A088999 001 Feb 05, 1991

50MG

A089000 001 Feb 05, 1991

75MG

A089001 001 Feb 05, 1991

PUREPAC PHARM

25MG

A089425 001 Jul 12, 1990

50MG

A089426 001 Jul 12, 1990

75MG

A089427 001 Jul 12, 1990

WATSON LABS

50MG

A087160 001 Jun 07, 1996

PERSANTINE

+ BOEHRINGER INGELHEIM

25MG

N012836 003 Dec 22, 1986

+

50MG

N012836 004 Feb 06, 1987

+

75MG

N012836 005 Feb 06, 1987

DIRITHROMYCIN

TABLET, DELAYED RELEASE; ORAL

DYNABAC

LILLY RES LABS

250MG

N050678 001 Jun 19, 1995

DISOPYRAMIDE PHOSPHATE

CAPSULE; ORAL

DISOPYRAMIDE PHOSPHATE

INTERPHARM

EQ 100MG BASE

A071190 001 Jan 15, 1987

EQ 150MG BASE

A071191 001 Jan 15, 1987

IVAX SUB TEVA PHARMS

EQ 100MG BASE

A070186 001 Nov 18, 1985

EQ 150MG BASE

A070187 001 Nov 18, 1985

MYLAN

EQ 100MG BASE

A070138 001 Jun 14, 1985

EQ 150MG BASE

A070139 001 Jun 14, 1985

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DISOPYRAMIDE PHOSPHATE

CAPSULE;ORAL

DISOPYRAMIDE PHOSPHATE

RISING	EQ 100MG BASE	A070470	001	Dec 10, 1985
	EQ 150MG BASE	A070471	001	Dec 10, 1985
SUN PHARM INDUSTRIES	EQ 100MG BASE	A070351	001	Dec 17, 1985
	EQ 150MG BASE	A070352	001	Dec 17, 1985
SUPERPHARM	EQ 100MG BASE	A070940	001	Feb 09, 1987
	EQ 150MG BASE	A070941	001	Feb 09, 1987
WATSON LABS	EQ 100MG BASE	A070240	001	Feb 02, 1986
	EQ 150MG BASE	A070241	001	Feb 02, 1986

CAPSULE, EXTENDED RELEASE;ORAL

DISOPYRAMIDE PHOSPHATE

NESHER PHARMS	EQ 150MG BASE	A071200	001	Dec 15, 1987
---------------	---------------	---------	-----	--------------

DISULFIRAM

TABLET;ORAL

ANTABUSE

+ ODYSSEY PHARMS	250MG	A088482	001	Dec 08, 1983
+	500MG	A088483	001	Dec 08, 1983
+ TEVA WOMENS	250MG **	N007883	003	
+	500MG **	N007883	002	

DISULFIRAM

DASH PHARMS	250MG	A203916	001	Mar 04, 2015
	500MG	A203916	002	Mar 04, 2015
HIKMA	250MG	A202652	001	Feb 05, 2014
	500MG	A202652	002	Feb 05, 2014
STRIDES PHARMA	250MG	A088792	001	Aug 14, 1984
	500MG	A088793	001	Aug 14, 1984
WATSON LABS	250MG	A086889	001	
	250MG	A087973	001	Aug 05, 1983
	500MG	A087974	001	Aug 05, 1983
WATSON LABS TEVA	500MG	A086890	001	

DIVALPROEX SODIUM

CAPSULE, DELAYED REL PELLETS;ORAL

DIVALPROEX SODIUM

RISING	EQ 125MG VALPROIC ACID	A090407	001	Mar 28, 2011
TEVA PHARMS USA	EQ 125MG VALPROIC ACID	A211505	001	Nov 17, 2020

TABLET, DELAYED RELEASE;ORAL

DEPAKOTE CP

ABBOTT	EQ 250MG BASE	N019794	001	Jul 11, 1990
	EQ 500MG BASE	N019794	002	Jul 11, 1990

DIVALPROEX SODIUM

ACTAVIS LABS FL INC	EQ 500MG VALPROIC ACID	A079080	001	Feb 25, 2011
ENDO OPERATIONS	EQ 500MG VALPROIC ACID	A078411	001	Nov 03, 2008
MYLAN	EQ 125MG VALPROIC ACID	A077254	001	Jul 29, 2008
	EQ 250MG VALPROIC ACID	A077254	002	Jul 29, 2008
	EQ 500MG VALPROIC ACID	A077254	003	Jul 29, 2008
NORVIUM BIOSCIENCE	EQ 125MG VALPROIC ACID	A090062	001	Mar 17, 2009
	EQ 250MG VALPROIC ACID	A090062	002	Mar 17, 2009
	EQ 500MG VALPROIC ACID	A090062	003	Mar 17, 2009
TEVA	EQ 125MG VALPROIC ACID	A076941	001	Jul 29, 2008
	EQ 250MG VALPROIC ACID	A076941	002	Jul 29, 2008
	EQ 500MG VALPROIC ACID	A076941	003	Jul 29, 2008

TABLET, EXTENDED RELEASE;ORAL

DIVALPROEX SODIUM

COSETTE	EQ 500MG VALPROIC ACID	A078700	001	Aug 03, 2009
ENDO OPERATIONS	EQ 250MG VALPROIC ACID	A078445	001	Feb 26, 2009
	EQ 500MG VALPROIC ACID	A078445	002	Aug 04, 2009
IMPAX LABS	EQ 250MG VALPROIC ACID	A078791	001	May 06, 2009
	EQ 500MG VALPROIC ACID	A078791	002	Aug 04, 2009
UTOPIC PHARMS	EQ 250MG VALPROIC ACID	A214462	001	Mar 15, 2021
	EQ 500MG VALPROIC ACID	A214462	002	Mar 15, 2021

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DOBUTAMINE HYDROCHLORIDE

INJECTABLE; INJECTION

DOBUTAMINE HYDROCHLORIDE

BAXTER HLTHCARE	EQ 12.5MG BASE/ML	A074381	001	Sep 26, 1996
DR REDDYS	EQ 12.5MG BASE/ML	A074995	001	Mar 31, 1998
HOSPIRA	EQ 12.5MG BASE/ML	A074292	001	Feb 16, 1995
	EQ 1.25GM BASE/100ML	A074634	001	Sep 27, 1996
LUITPOLD	EQ 12.5MG BASE/ML	A074545	001	Jun 25, 1998
TELIGENT	EQ 12.5MG BASE/ML	A074098	001	Feb 21, 1995
TEVA PARENTERAL	EQ 12.5MG BASE/ML	A074206	001	Oct 19, 1993
WATSON LABS	EQ 12.5MG BASE/ML	A074114	001	Nov 30, 1993
WATSON LABS INC	EQ 12.5MG BASE/ML	A074279	001	Feb 18, 1998
DOBUTAMINE HYDROCHLORIDE IN	DEXTROSE 5%			
HOSPIRA	EQ 50MG BASE/100ML	N020269	001	Oct 19, 1993
	EQ 100MG BASE/100ML	N020269	002	Oct 19, 1993
	EQ 200MG BASE/100ML	N020269	003	Oct 19, 1993
DOBUTREX				
+ LILLY	EQ 12.5MG BASE/ML **	N017820	002	

DOCETAXEL

INJECTABLE; INJECTION

DOCEFREZ

+ SUN PHARM	20MG/VIAL	N022534	001	May 03, 2011
	80MG/VIAL	N022534	002	May 03, 2011

DOCETAXEL

+ ACCORD HLTHCARE	20MG/0.5ML (40MG/ML)	N201195	001	Jun 08, 2011
	80MG/2ML (40MG/ML)	N201195	002	Jun 08, 2011
+ APOTEX INC	20MG/0.5ML (40MG/ML)	N022312	001	Jan 11, 2012
	80MG/2ML (40MG/ML)	N022312	002	Jan 11, 2012
DFB ONCOLOGY LTD	20MG/ML (20MG/ML)	A206177	001	Jan 20, 2017
	80MG/4ML (20MG/ML)	A206177	002	Jan 20, 2017
	200MG/10ML (20MG/ML)	A206177	003	Jan 20, 2017
HENGRUI PHARMA	40MG/ML	A203170	001	Feb 15, 2017
+ HOSPIRA INC	120MG/6ML (20MG/ML)	N022234	006	Jun 23, 2016
NORVIUM BIOSCIENCE	20MG/ML (20MG/ML)	A203892	001	May 11, 2023
	80MG/4ML (20MG/ML)	A203892	002	May 11, 2023
NOVAST LABS	20MG/2ML (10MG/ML)	A207563	001	Aug 31, 2017
PFIZER LABS	20MG/2ML (10MG/ML)	N202356	001	Mar 13, 2014
	80MG/8ML (10MG/ML)	N202356	002	Mar 13, 2014
	130MG/13ML (10MG/ML)	N202356	003	Mar 13, 2014
	200MG/20ML (10MG/ML)	N202356	004	Mar 13, 2014
SUN PHARM	20MG/ML (20MG/ML)	N022534	003	Jan 08, 2019
	80MG/4ML (20MG/ML)	N022534	004	Jan 08, 2019
	160MG/8ML (20MG/ML)	N022534	005	Jan 08, 2019
TEVA PHARMS USA	20MG/ML (20MG/ML)	A203877	001	Sep 16, 2015
	80MG/4ML (20MG/ML)	A203877	002	Sep 16, 2015
TAXOTERE				
+ SANOFI AVENTIS US	40MG/ML **	N020449	001	May 14, 1996

DOFETILIDE

CAPSULE; ORAL

DOFETILIDE

NOVAST LABS	0.125MG	A212410	001	Dec 27, 2019
	0.25MG	A212410	002	Dec 27, 2019
	0.5MG	A212410	003	Dec 27, 2019
PRINSTON INC	0.125MG	A211223	001	Dec 17, 2019
	0.25MG	A211223	002	Dec 17, 2019
	0.5MG	A211223	003	Dec 17, 2019
RK PHARMA	0.125MG	A215323	001	Apr 14, 2022
	0.25MG	A215323	002	Apr 14, 2022
	0.5MG	A215323	003	Apr 14, 2022
TEVA PHARMS USA	0.125MG	A210018	001	Apr 15, 2022
	0.25MG	A210018	002	Apr 15, 2022
	0.5MG	A210018	003	Apr 15, 2022

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DOLASETRON MESYLATE

INJECTABLE; INJECTION

ANZEMET

+	VALIDUS PHARMS	12.5MG/0.625ML (20MG/ML)	N020624	002	Sep 11, 1997
+		100MG/5ML (20MG/ML)	N020624	001	Sep 11, 1997
		500MG/25ML (20MG/ML)	N020624	003	Dec 11, 2001

TABLET; ORAL

ANZEMET

+	VALIDUS PHARMS	50MG	N020623	001	Sep 11, 1997
+		100MG	N020623	002	Sep 11, 1997

DOLUTEGRAVIR SODIUM

TABLET; ORAL

TIVICAY

+	VIIV HLTHCARE	EQ 10MG BASE **	N204790	002	Jun 09, 2016
+		EQ 25MG BASE **	N204790	003	Jun 09, 2016

DONEPEZIL HYDROCHLORIDE

SOLUTION; ORAL

ARICEPT

+	EISAI INC	5MG/5ML	N021719	001	Oct 18, 2004
---	-----------	---------	---------	-----	--------------

TABLET; ORAL

DONEPEZIL HYDROCHLORIDE

	ACCORD HLTHCARE	5MG	A201335	001	Aug 29, 2011
		10MG	A201335	002	Aug 29, 2011
	ACTAVIS ELIZABETH	23MG	A202415	001	Dec 17, 2015
	APOTEX	5MG	A078841	001	Jun 02, 2011
		10MG	A078841	002	Jun 02, 2011
	CHARTWELL RX	23MG	A203419	001	Apr 12, 2016
	ENDO OPERATIONS	23MG	A202542	001	Jul 24, 2013
	HERITAGE PHARMA	5MG	A077344	001	May 31, 2011
		10MG	A077344	002	May 31, 2011
	HIKMA PHARMS	5MG	A090247	001	May 31, 2011
		10MG	A090247	002	May 31, 2011
	HISUN PHARM HANGZHOU	23MG	A202410	001	Mar 24, 2017
	NATCO PHARMA	5MG	A090521	001	May 31, 2011
		10MG	A090521	002	May 31, 2011
	OSMOTICA PHARM US	23MG	A203114	001	Jan 26, 2016
	RISING	23MG	A202656	001	Oct 22, 2015
	SANDOZ	5MG	A090290	001	May 31, 2011
		10MG	A090290	002	May 31, 2011
	SUN PHARM	23MG	A204293	001	Jun 05, 2015
	SUN PHARM INDS	5MG	A090493	001	May 31, 2011
		10MG	A090493	002	May 31, 2011
	SUN PHARM INDS LTD	5MG	A076786	001	Nov 26, 2010
		10MG	A076786	002	Nov 26, 2010
	UNICHEM	5MG	A203656	001	Jun 23, 2016
		10MG	A203656	002	Jun 23, 2016
	WOCKHARDT	5MG	A091267	001	May 31, 2011
		10MG	A091267	002	May 31, 2011

TABLET, ORALLY DISINTEGRATING; ORAL

ARICEPT ODT

+	EISAI INC	5MG	N021720	001	Oct 18, 2004
+		10MG	N021720	002	Oct 18, 2004

DONEPEZIL HYDROCHLORIDE

	CHARTWELL RX	5MG	A078388	002	Nov 26, 2010
		10MG	A078388	001	Nov 26, 2010
	SANDOZ	5MG	A091198	001	May 10, 2011
		10MG	A091198	002	May 10, 2011
	SUN PHARM INDUSTRIES	5MG	A077975	002	Dec 11, 2009
		10MG	A077975	001	Dec 11, 2009
	UNICHEM	5MG	A204831	001	Nov 10, 2016
		10MG	A204831	002	Nov 10, 2016
	ZYDUS PHARMS USA INC	5MG	A090175	001	May 10, 2011
		10MG	A090175	002	May 10, 2011

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DOPAMINE HYDROCHLORIDE

INJECTABLE; INJECTION

DOPAMINE HYDROCHLORIDE

ABBOTT	40MG/ML	A070656	001	Jan 24, 1989
	80MG/ML	A070657	001	Jan 24, 1989
ABRAXIS PHARM	40MG/ML	A070012	001	Jun 12, 1985
	40MG/ML	A070058	001	Mar 20, 1985
	80MG/ML	A070013	001	Jun 12, 1985
	80MG/ML	A070059	001	Mar 20, 1985
	160MG/ML	A070364	001	Dec 04, 1985
AM REGENT	40MG/ML	A070799	001	Feb 11, 1987
	80MG/ML	A070820	001	Feb 11, 1987
	160MG/ML	A070826	001	Feb 11, 1987
BAXTER HLTHCARE	40MG/ML	N018398	001	
	80MG/ML	N018398	002	Mar 22, 1982
HOSPIRA	40MG/ML	A074403	001	May 23, 1996
IGI LABS INC	40MG/ML	A070087	001	Oct 23, 1985
	80MG/ML	A070089	001	Oct 23, 1985
	80MG/ML	A070090	001	Oct 23, 1985
	80MG/ML	A070091	001	Oct 23, 1985
	160MG/ML	A070092	001	Oct 23, 1985
	160MG/ML	A070093	001	Oct 23, 1985
	160MG/ML	A070094	001	Oct 23, 1985
INTL MEDICATION	40MG/ML	N018014	001	
LYPHOMED	40MG/ML	N018549	001	Mar 11, 1983
SMITH AND NEPHEW	40MG/ML	A070011	001	Aug 29, 1985
	40MG/ML	A070046	001	Aug 29, 1985
	80MG/ML	A070047	001	Aug 29, 1985
TELIGENT	40MG/ML	N018656	001	Jun 28, 1983
TEVA PARENTERAL	40MG/ML	A072999	001	Oct 23, 1991
	80MG/ML	A073000	001	Oct 23, 1991
WARNER CHILCOTT	40MG/ML	A070558	001	Sep 20, 1985
	40MG/ML	N018138	001	
	80MG/ML	A070559	001	Sep 20, 1985
DOPAMINE HYDROCHLORIDE AND DEXTROSE 5%				
+ B BRAUN	80MG/100ML	N019099	002	Oct 15, 1986
+	320MG/100ML	N019099	004	Oct 15, 1986
DOPAMINE HYDROCHLORIDE AND DEXTROSE 5% IN PLASTIC CONTAINER				
+ B BRAUN	40MG/100ML	N019099	001	Oct 15, 1986
+	160MG/100ML	N019099	003	Oct 15, 1986
DOPAMINE HYDROCHLORIDE IN DEXTROSE 5%				
HOSPIRA	1.6MG/ML	N020542	001	Aug 30, 1995
INTROPIN				
HOSPIRA	40MG/ML	N017395	001	
	80MG/ML	N017395	002	
	160MG/ML	N017395	003	

DORIPENEM

INJECTABLE; INTRAVENOUS

DORIBAX

+ SHIONOGI INC	250MG/VIAL **	N022106	002	Oct 05, 2010
+	500MG/VIAL **	N022106	001	Oct 12, 2007

DORZOLAMIDE HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

DORZOLAMIDE HYDROCHLORIDE

AM REGENT	EQ 2% BASE	A079186	001	Nov 18, 2009
HIKMA	EQ 2% BASE	A077846	001	Oct 28, 2008
TEVA PHARMS	EQ 2% BASE	A078756	001	Dec 04, 2008
ZAMBON SPA	EQ 2% BASE	A091034	001	Dec 04, 2013
TRUSOPT				
+ MSD SUB MERCK	EQ 2% BASE **	N020408	001	Dec 09, 1994

DORZOLAMIDE HYDROCHLORIDE; TIMOLOL MALEATE

SOLUTION/DROPS; OPHTHALMIC

DORZOLAMIDE HYDROCHLORIDE AND TIMOLOL MALEATE

CHARTWELL MOLECULAR	EQ 2% BASE;EQ 0.5% BASE	A201998	001	Dec 17, 2014
HIKMA	EQ 2% BASE;EQ 0.5% BASE	A077847	001	Oct 28, 2008
RUBICON	EQ 2% BASE;EQ 0.5% BASE	A078201	001	Oct 28, 2008
TEVA PHARMS	EQ 2% BASE;EQ 0.5% BASE	A078704	001	Sep 28, 2009
ZAMBON SPA	EQ 2% BASE;EQ 0.5% BASE	A091180	001	Dec 04, 2013

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DOXACURIUM CHLORIDEINJECTABLE; INJECTION
NUROMAX

ABBVIE	EQ 1MG BASE/ML	N019946 001	Mar 07, 1991
--------	----------------	-------------	--------------

DOXAPRAM HYDROCHLORIDEINJECTABLE; INJECTION
DOXAPRAM HYDROCHLORIDE

WATSON LABS	20MG/ML	A073529 001	Jan 30, 1992
-------------	---------	-------------	--------------

DOXAZOSIN MESYLATETABLET; ORAL
DOXAZOSIN MESYLATE

ACTAVIS ELIZABETH	EQ 1MG BASE	A075574 001	Oct 18, 2000
	EQ 2MG BASE	A075574 002	Oct 18, 2000
	EQ 4MG BASE	A075574 003	Oct 18, 2000
	EQ 8MG BASE	A075574 004	Oct 18, 2000
ANI PHARMS	EQ 1MG BASE	A075432 001	Oct 18, 2000
	EQ 2MG BASE	A075432 002	Oct 18, 2000
	EQ 4MG BASE	A075432 003	Oct 18, 2000
	EQ 8MG BASE	A075432 004	Oct 18, 2000
CHARTWELL RX	EQ 1MG BASE	A075646 001	Oct 18, 2000
	EQ 2MG BASE	A075646 002	Oct 18, 2000
	EQ 4MG BASE	A075646 003	Oct 18, 2000
	EQ 8MG BASE	A075646 004	Oct 18, 2000
GENPHARM	EQ 1MG BASE	A075466 001	Oct 18, 2000
	EQ 2MG BASE	A075466 002	Oct 18, 2000
	EQ 4MG BASE	A075466 003	Oct 18, 2000
	EQ 8MG BASE	A075466 004	Oct 18, 2000
IVAX SUB TEVA PHARMS	EQ 1MG BASE	A075453 001	Oct 18, 2000
	EQ 2MG BASE	A075453 002	Oct 18, 2000
	EQ 4MG BASE	A075453 003	Oct 18, 2000
	EQ 8MG BASE	A075453 004	Oct 18, 2000
NESHER PHARMS	EQ 1MG BASE	A075609 001	Oct 18, 2000
	EQ 2MG BASE	A075609 002	Oct 18, 2000
	EQ 4MG BASE	A075609 003	Oct 18, 2000
	EQ 8MG BASE	A075609 004	Oct 18, 2000
PLIVA	EQ 1MG BASE	A075750 001	Jun 08, 2001
	EQ 2MG BASE	A075750 002	Jun 08, 2001
	EQ 4MG BASE	A075750 003	Jun 08, 2001
	EQ 8MG BASE	A075750 004	Jun 08, 2001
STRIDES PHARMA	EQ 1MG BASE	A076161 001	Jun 10, 2004
	EQ 2MG BASE	A076161 002	Jun 10, 2004
	EQ 4MG BASE	A076161 003	Jun 10, 2004
	EQ 8MG BASE	A076161 004	Jun 10, 2004
TEVA	EQ 1MG BASE	A075353 001	Jan 12, 2001
	EQ 2MG BASE	A075353 002	Jan 12, 2001
	EQ 4MG BASE	A075353 003	Jan 12, 2001
	EQ 8MG BASE	A075353 004	Jan 12, 2001
WATSON LABS INC	EQ 1MG BASE	A075426 001	Oct 18, 2000
	EQ 2MG BASE	A075426 002	Oct 18, 2000
	EQ 4MG BASE	A075426 003	Oct 18, 2000
	EQ 8MG BASE	A075426 004	Oct 18, 2000

DOXEPIN HYDROCHLORIDECAPSULE; ORAL
DOXEPIN HYDROCHLORIDE

DAVA PHARMS INC	EQ 10MG BASE	A071685 001	Jan 05, 1988
	EQ 25MG BASE	A071686 001	Jan 05, 1988
	EQ 50MG BASE	A071673 001	Jan 05, 1988
	EQ 75MG BASE	A071674 001	Jan 05, 1988
	EQ 100MG BASE	A071675 001	Jan 05, 1988
ENDO OPERATIONS	EQ 150MG BASE	A071676 001	Jan 05, 1988
	EQ 10MG BASE	A071422 002	Nov 09, 1987
	EQ 25MG BASE	A071422 003	Nov 09, 1987
	EQ 50MG BASE	A071422 004	Nov 09, 1987
	EQ 75MG BASE	A071422 005	Nov 09, 1987
LANNETT CO INC	EQ 100MG BASE	A071422 001	Nov 09, 1987
	EQ 10MG BASE	A212997 001	Jul 24, 2020
	EQ 25MG BASE	A212997 002	Jul 24, 2020
	EQ 50MG BASE	A212997 003	Jul 24, 2020

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DOXEPIN HYDROCHLORIDE

CAPSULE; ORAL

DOXEPIN HYDROCHLORIDE

	EQ 75MG BASE	A212997 004	Jul 24, 2020
	EQ 100MG BASE	A212997 005	Jul 24, 2020
NEW RIVER	EQ 10MG BASE	N016987 001	
	EQ 25MG BASE	N016987 002	
	EQ 50MG BASE	N016987 003	
	EQ 75MG BASE	N016987 006	
	EQ 100MG BASE	N016987 004	
	EQ 150MG BASE	N016987 007	Apr 13, 1987
PUREPAC PHARM	EQ 10MG BASE	A073054 001	Dec 28, 1990
	EQ 25MG BASE	A072109 001	Dec 28, 1990
	EQ 50MG BASE	A073055 001	Dec 28, 1990
	EQ 75MG BASE	A072386 001	Sep 08, 1988
	EQ 100MG BASE	A072110 001	Sep 08, 1988
	EQ 150MG BASE	A072387 001	Sep 08, 1988
QUANTUM PHARMICS	EQ 10MG BASE	A070972 001	Sep 29, 1987
	EQ 25MG BASE	A070973 001	Sep 29, 1987
	EQ 50MG BASE	A070931 001	Sep 29, 1987
	EQ 75MG BASE	A070932 001	Sep 29, 1987
	EQ 100MG BASE	A072375 001	Mar 15, 1989
	EQ 150MG BASE	A072376 001	Mar 15, 1989
SANDOZ	EQ 10MG BASE	A071487 001	Mar 02, 1987
	EQ 25MG BASE	A070827 001	May 15, 1986
	EQ 50MG BASE	A070828 001	May 15, 1986
	EQ 75MG BASE	A070825 001	May 15, 1986
	EQ 100MG BASE	A071562 001	Mar 02, 1987
SUN PHARM INDUSTRIES	EQ 25MG BASE	A071502 001	Feb 18, 1988
	EQ 50MG BASE	A071653 001	Feb 18, 1988
	EQ 75MG BASE	A071654 001	Feb 18, 1988
	EQ 100MG BASE	A071521 001	Feb 18, 1988
WATSON LABS	EQ 10MG BASE	A070952 001	Mar 04, 1987
	EQ 10MG BASE	A071485 001	Apr 30, 1987
	EQ 10MG BASE	A072985 001	Mar 29, 1991
	EQ 25MG BASE	A070953 001	May 15, 1986
	EQ 25MG BASE	A071486 001	Apr 30, 1987
	EQ 25MG BASE	A072986 001	Mar 29, 1991
	EQ 50MG BASE	A070954 001	May 15, 1986
	EQ 50MG BASE	A071238 001	Apr 30, 1987
	EQ 75MG BASE	A071326 001	Apr 30, 1987
	EQ 75MG BASE	A071763 001	Feb 09, 1988
	EQ 100MG BASE	A070955 001	May 15, 1986
	EQ 100MG BASE	A071239 001	Apr 30, 1987
	EQ 150MG BASE	A071764 001	Feb 09, 1988
WATSON LABS TEVA	EQ 50MG BASE	A072987 001	Mar 29, 1991
SINEQUAN			
+ PFIZER	EQ 10MG BASE **	N016798 003	
+	EQ 25MG BASE **	N016798 001	
+	EQ 50MG BASE **	N016798 002	
+	EQ 75MG BASE **	N016798 006	
+	EQ 100MG BASE **	N016798 005	
+	EQ 150MG BASE **	N016798 007	
CONCENTRATE; ORAL			
DOXEPIN HYDROCHLORIDE			
PHARM ASSOC	EQ 10MG BASE/ML	A075924 001	Jan 15, 2004
PHARMOBEDIANT CNSLTG	EQ 10MG BASE/ML	A071918 001	Jul 20, 1988
TEVA PHARMS	EQ 10MG BASE/ML	A071609 001	Nov 09, 1987
SINEQUAN			
+ PFIZER	EQ 10MG BASE/ML **	N017516 001	
TABLET; ORAL			
DOXEPIN HYDROCHLORIDE			
AJANTA PHARMA LTD	EQ 3MG BASE	A218564 001	Jul 01, 2024
	EQ 6MG BASE	A218564 002	Jul 01, 2024
AUROBINDO PHARMA LTD	EQ 3MG BASE	A216041 001	Jul 25, 2024
	EQ 6MG BASE	A216041 002	Jul 25, 2024

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DOXERCALCIFEROL

CAPSULE; ORAL

DOXERCALCIFEROL

HIKMA

0.5MCG

A091433 001 Sep 23, 2011

1MCG

A091433 002 Jan 14, 2014

2.5MCG

A091433 003 Jan 14, 2014

INJECTABLE; INJECTION

DOXERCALCIFEROL

AMNEAL

2MCG/ML (2MCG/ML)

A208974 001 May 24, 2017

4MCG/2ML (2MCG/ML)

A208974 002 May 24, 2017

4MCG/2ML (2MCG/ML)

A208975 001 May 24, 2017

EPIC PHARMA LLC

2MCG/ML (2MCG/ML)

A203929 002 Mar 28, 2016

4MCG/2ML (2MCG/ML)

A203929 001 May 07, 2015

+ HOSPIRA

10MCG/5ML (2MCG/ML)

N208614 002 Jul 24, 2018

SUN PHARM

2MCG/ML (2MCG/ML)

A203875 001 Nov 14, 2019

4MCG/2ML (2MCG/ML)

A203875 002 Nov 14, 2019

DOXORUBICIN HYDROCHLORIDE

INJECTABLE; INJECTION

ADRIAMYCIN PFS

PFIZER

2MG/ML

A063165 001 Jan 30, 1991

200MG/100ML

A063165 002 Jan 30, 1991

DOXORUBICIN HYDROCHLORIDE

ALMAJECT

2MG/ML

A065515 001 Nov 08, 2012

HISUN PHARM HANGZHOU

20MG/VIAL

A206062 001 May 13, 2019

HLTHCARE

2MG/ML

A200146 001 Jul 18, 2012

MYLAN LABS LTD

10MG/VIAL

A200170 001 Oct 28, 2011

NORVIUM BIOSCIENCE

2MG/ML

A200901 001 Feb 14, 2012

PFIZER

10MG/VIAL

N050467 001

20MG/VIAL

N050467 003 May 20, 1985

50MG/VIAL

N050467 002

150MG/VIAL

N050467 004 Jul 22, 1987

+ PHARMACHEMIE BV

2MG/ML

N050629 003 Mar 28, 2011

10MG/VIAL

A063336 001 Feb 28, 1995

20MG/VIAL

A063097 001 May 21, 1990

50MG/VIAL

A063097 002 May 21, 1990

200MG/100ML

A063097 003 May 21, 1990

TEVA PHARMS USA

2MG/ML

A063336 004 Feb 28, 1995

200MG/100ML

A064140 001 Jul 28, 1995

A064140 002 Jul 28, 1995

RUBEX

BRISTOL MYERS SQUIBB

10MG/VIAL

A062926 001 Apr 13, 1989

50MG/VIAL

A062926 002 Apr 13, 1989

100MG/VIAL

A062926 003 Apr 13, 1989

DOXYCYCLINE

CAPSULE; ORAL

DOXYCYCLINE

COSETTE

EQ 50MG BASE

A204446 001 May 28, 2015

EQ 75MG BASE

A204446 002 May 28, 2015

EQ 100MG BASE

A204446 003 May 28, 2015

IMPAX LABS INC

EQ 150MG BASE

A200065 001 Feb 17, 2011

RISING

EQ 50MG BASE

A208942 001 Jan 21, 2021

EQ 75MG BASE

A208942 002 Jan 21, 2021

EQ 100MG BASE

A208942 003 Jan 21, 2021

SANDOZ INC

EQ 50MG BASE

A065032 001 Jun 30, 2000

EQ 100MG BASE

A065032 002 Jun 30, 2000

STRIDES PHARMA

EQ 75MG BASE

A065055 004 Apr 18, 2005

EQ 150MG BASE

A202778 001 Jun 08, 2012

WATSON LABS

EQ 50MG BASE

A065041 001 Apr 28, 2000

EQ 100MG BASE

A065041 002 Apr 28, 2000

FOR SUSPENSION; ORAL

DOXYCHEL

RACHELLE

EQ 25MG BASE/5ML

A061720 001

VIBRAMYCIN

+ PFIZER

EQ 25MG BASE/5ML

N050006 001

TABLET; ORAL

DOXYCYCLINE

SANDOZ INC

EQ 50MG BASE

A065353 001 Nov 27, 2006

EQ 75MG BASE

A065353 002 Nov 27, 2006

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DOXYCYCLINE

TABLET;ORAL

DOXYCYCLINE

	EQ 100MG BASE	A065353 003	Nov 27, 2006
SOMERSET THERAPS LLC	EQ 50MG BASE	A065377 001	Nov 07, 2006
	EQ 75MG BASE	A065377 002	Nov 07, 2006
	EQ 100MG BASE	A065377 003	Nov 07, 2006
	EQ 150MG BASE	A065427 001	Jun 07, 2007
SUN PHARM INDUSTRIES	EQ 50MG BASE	A065471 001	Apr 17, 2009
	EQ 75MG BASE	A065471 002	Apr 17, 2009
	EQ 100MG BASE	A065471 003	Apr 17, 2009

DOXYCYCLINE CALCIUM

SUSPENSION;ORAL

VIBRAMYCIN

+ PFIZER

EQ 50MG BASE/5ML

N050480 001

DOXYCYCLINE HYCLATE

CAPSULE;ORAL

ACTICLATE CAP

+ CHARTWELL RX

EQ 75MG BASE

N208253 001 Apr 26, 2016

DOXY-LEMMON

TEVA

EQ 50MG BASE

A062497 001 Aug 23, 1984

EQ 100MG BASE

A062497 002 Jun 15, 1984

DOXYCYCLINE HYCLATE

AJANTA PHARMA LTD

EQ 50MG BASE

A211012 001 Sep 24, 2018

EQ 100MG BASE

A211012 002 Sep 24, 2018

HALSEY

EQ 50MG BASE

A062119 002 May 24, 1985

EQ 100MG BASE

A062119 001 May 24, 1985

HEATHER

EQ 50MG BASE

A062463 001 Dec 07, 1983

EQ 100MG BASE

A062463 002 Dec 07, 1983

HIKMA INTL PHARMS

EQ 20MG BASE

A065103 001 May 13, 2005

INTERPHARM

EQ 50MG BASE

A062763 001 Sep 02, 1988

EQ 100MG BASE

A062763 002 Sep 02, 1988

MUTUAL PHARM

EQ 50MG BASE

A062418 001 Jan 28, 1983

EQ 100MG BASE

A062418 002 Jan 28, 1983

NESHER PHARMS

EQ 50MG BASE

A208263 001 Nov 22, 2021

EQ 100MG BASE

A208263 002 Nov 22, 2021

NOSTRUM LABS INC

EQ 50MG BASE

A209393 001 Dec 10, 2020

EQ 100MG BASE

A209393 002 Dec 10, 2020

PAR PHARM

EQ 50MG BASE

A062434 001 Oct 19, 1984

EQ 100MG BASE

A062442 001 Dec 22, 1983

PVT FORM

EQ 50MG BASE

A062631 001 Jul 24, 1986

EQ 100MG BASE

A062631 002 Jul 24, 1986

RANBAXY

EQ 50MG BASE

A062479 001 Dec 23, 1983

EQ 100MG BASE

A062479 002 Dec 23, 1983

STRIDES PHARMA

EQ 50MG BASE

A062337 001 Mar 29, 1982

EQ 100MG BASE

A062337 002 Mar 29, 1982

SUPERPHARM

EQ 50MG BASE

A062469 001 Oct 31, 1984

EQ 100MG BASE

A062469 002 Oct 31, 1984

WARNER CHILCOTT

EQ 50MG BASE

A062594 001 Dec 05, 1985

EQ 100MG BASE

A062594 002 Dec 05, 1985

WATSON LABS

EQ 50MG BASE

A061717 001

EQ 50MG BASE

A062142 001

EQ 100MG BASE

A061717 002

EQ 100MG BASE

A062142 002

ZHEJIANG YONGTAI

EQ 50MG BASE

A212610 001 Mar 31, 2020

EQ 100MG BASE

A212610 002 Mar 31, 2020

PERIOSTAT

+ COLLAGENEX

EQ 20MG BASE **

N050744 001 Sep 30, 1998

VIBRAMYCIN

+ PFIZER

EQ 50MG BASE **

N050007 001

CAPSULE, COATED PELLETS;ORAL

DOXYCYCLINE HYCLATE

PLIVA

EQ 100MG BASE

A063187 001 Jun 30, 1992

CAPSULE, DELAYED RELEASE;ORAL

DORYX

+ MAYNE PHARMA

EQ 75MG BASE

N050582 002 Aug 13, 2001

+

EQ 100MG BASE

N050582 001 Jul 22, 1985

WARNER CHILCOTT

EQ 100MG BASE

A062653 001 Oct 30, 1985

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DOXYCYCLINE HYCLATE

CAPSULE, DELAYED RELEASE;ORAL

DOXYCYCLINE HYCLATE

BAUSCH

EQ 75MG BASE

A065281 001 Dec 21, 2005

EQ 100MG BASE

A065281 002 Dec 21, 2005

INJECTABLE; INJECTION

DOXYCHEL HYCLATE

RACHELLE

EQ 100MG BASE/VIAL

A061953 001

DOXYCYCLINE

HIKMA

EQ 100MG BASE/VIAL

A062450 001 Oct 27, 1983

EQ 100MG BASE/VIAL

A062569 001 Mar 09, 1988

EQ 200MG BASE/VIAL

A062450 002 Oct 27, 1983

EQ 200MG BASE/VIAL

A062569 002 Mar 09, 1988

VIBRAMYCIN

+ PFIZER

EQ 100MG BASE/VIAL **

N050442 002

+

EQ 200MG BASE/VIAL **

N050442 001

SYSTEM, EXTENDED RELEASE;PERIODONTAL

ATRIDOX

+ DEN-MAT

50MG

N050751 001 Sep 03, 1998

TABLET; ORAL

DOXY-LEMMON

TEVA

EQ 100MG BASE

A062581 001 Mar 15, 1985

DOXYCYCLINE HYCLATE

ALEMBIC

EQ 100MG BASE

A210536 001 May 14, 2020

AMNEAL

EQ 100MG BASE

A216599 001 Oct 24, 2022

AMNEAL PHARMS CO

EQ 75MG BASE

A209372 001 Oct 06, 2017

EQ 150MG BASE

A209372 002 Oct 06, 2017

AVET LIFESCIENCES

EQ 100MG BASE

A209969 001 Nov 09, 2018

HEATHER

EQ 100MG BASE

A062462 001 May 11, 1983

HERITAGE PHARMA

EQ 20MG BASE

A065163 001 May 13, 2005

INTERPHARM

EQ 100MG BASE

A062764 001 Sep 02, 1988

MUTUAL PHARM

EQ 100MG BASE

A062391 001 Sep 30, 1982

NOSTRUM LABS INC

EQ 100MG BASE

A209560 001 Apr 15, 2024

ORYZA

EQ 100MG BASE

A213475 001 Mar 10, 2021

PRAXGEN

EQ 100MG BASE

A212487 001 Mar 30, 2022

RISING

EQ 75MG BASE

A209987 001 Oct 05, 2020

EQ 150MG BASE

A209987 002 Oct 05, 2020

STRIDES PHARMA

EQ 100MG BASE

A062538 001 Apr 07, 1986

SUPERPHARM

EQ 100MG BASE

A062494 001 Feb 20, 1985

TORRENT

EQ 75MG BASE

A214356 001 Apr 15, 2024

EQ 150MG BASE

A214356 002 Apr 15, 2024

WARNER CHILCOTT

EQ 100MG BASE

A062593 001 Aug 28, 1985

WATSON LABS

EQ 50MG BASE

A062392 001 Mar 31, 1983

EQ 100MG BASE

A062392 002 Mar 31, 1983

LYMEPAK

+ CHARTWELL PHARMA

EQ 100MG BASE

N209844 001 Jun 15, 2018

PERIOSTAT

+ GALDERMA LABS LP

EQ 20MG BASE **

N050783 001 Feb 02, 2001

VIBRA-TABS

+ PFIZER

EQ 100MG BASE **

N050533 001

TABLET, DELAYED RELEASE;ORAL

DORYX

+ MAYNE PHARMA

EQ 50MG BASE **

N050795 006 Dec 19, 2014

+

EQ 75MG BASE **

N050795 001 May 06, 2005

+

EQ 100MG BASE **

N050795 002 May 06, 2005

+

EQ 150MG BASE **

N050795 003 Jun 20, 2008

DORYX MPC

+ MAYNE PHARMA

EQ 120MG BASE **

N050795 008 May 20, 2016

DOXYCYCLINE HYCLATE

AUROBINDO PHARMA USA

EQ 150MG BASE

A091052 001 Feb 08, 2012

IMPAX LABS INC

EQ 75MG BASE

A090505 001 Dec 28, 2010

EQ 100MG BASE

A090505 002 Dec 28, 2010

LUPIN

EQ 50MG BASE

A208741 001 Aug 11, 2023

EQ 60MG BASE

A208741 002 Aug 11, 2023

EQ 75MG BASE

A208741 003 Aug 11, 2023

EQ 80MG BASE

A208741 004 Aug 11, 2023

EQ 100MG BASE

A208741 005 Aug 11, 2023

EQ 120MG BASE

A208741 006 Aug 11, 2023

EQ 150MG BASE

A208741 007 Aug 11, 2023

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DOXYCYCLINE HYCLATETABLET, DELAYED RELEASE;ORAL
DOXYCYCLINE HYCLATE

	EQ 200MG BASE	A208741 008	Aug 11, 2023
RISING	EQ 50MG BASE	A090431 003	May 23, 2016
	EQ 75MG BASE	A090431 001	Dec 28, 2010
	EQ 80MG BASE	A090431 004	Apr 29, 2016
	EQ 100MG BASE	A090431 002	Dec 28, 2010
	EQ 200MG BASE	A090431 005	May 19, 2016
ZYDUS PHARMS	EQ 75MG BASE	A206772 001	Dec 21, 2018
	EQ 100MG BASE	A206772 002	Dec 21, 2018
	EQ 150MG BASE	A206772 003	Dec 21, 2018

DOXYLAMINE SUCCINATE

CAPSULE;ORAL

UNISOM

PFIZER	25MG	N019440 001	Feb 05, 1986
--------	------	-------------	--------------

TABLET;ORAL

DECAPRYN

SANOFI AVENTIS US	12.5MG	N006412 015	
	25MG	N006412 014	

DOXY-SLEEP-AID

PAR PHARM	25MG	A070156 001	Jul 02, 1987
-----------	------	-------------	--------------

DOXYLAMINE SUCCINATE

COPLY PHARM	25MG	A088900 002	Feb 12, 1988
QUANTUM PHARMICS	25MG	A088603 001	Aug 07, 1984

DOXYLAMINE SUCCINATE; PYRIDOXINE HYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL

BENDECTIN

SANOFI AVENTIS US	10MG;10MG **	N010598 002	
-------------------	--------------	-------------	--

DOXYLAMINE SUCCINATE AND PYRIDOXINE HYDROCHLORIDE

ACTAVIS LABS FL INC	20MG;20MG	A212472 001	Mar 01, 2022
---------------------	-----------	-------------	--------------

DROMOSTANOLONE PROPIONATE

INJECTABLE;INJECTION

DROLBAN

LILLY	50MG/ML	N012936 001	
-------	---------	-------------	--

DRONABINOL

CAPSULE;ORAL

DRONABINOL

HIKMA	2.5MG	A079217 001	Jun 20, 2014
	5MG	A079217 002	Jun 20, 2014
	10MG	A079217 003	Jun 20, 2014
INSYS THERAP	2.5MG	A078501 001	Aug 19, 2011
	5MG	A078501 002	Aug 19, 2011
	10MG	A078501 003	Aug 19, 2011
LANNETT CO INC	2.5MG	A201463 001	May 18, 2018
	5MG	A201463 002	May 18, 2018
	10MG	A201463 003	May 18, 2018
SVC PHARMA	2.5MG	A078292 001	Jun 27, 2008
	5MG	A078292 002	Jun 27, 2008
	10MG	A078292 003	Jun 27, 2008

SOLUTION;ORAL

SYNDROS

+ BENUVIA OPERATIONS	5MG/ML	N205525 001	Mar 23, 2017
----------------------	--------	-------------	--------------

DROPERIDOL

INJECTABLE;INJECTION

DROPERIDOL

ABRAXIS PHARM	2.5MG/ML	A070992 001	Nov 17, 1986
	2.5MG/ML	A070993 001	Nov 17, 1986
ASTRAZENECA	2.5MG/ML	A072018 001	Oct 20, 1988
HOSPIRA	2.5MG/ML	A071645 001	Apr 07, 1988
	2.5MG/ML	A071981 001	Feb 29, 1988
	2.5MG/ML	A072272 001	Aug 31, 1995
IGI LABS INC	2.5MG/ML	A072019 001	Oct 19, 1988
	2.5MG/ML	A072020 001	Oct 19, 1988
	2.5MG/ML	A072021 001	Oct 19, 1988
LUITPOLD	2.5MG/ML	A072335 001	Oct 24, 1988
SMITH AND NEPHEW	2.5MG/ML	A071750 001	Sep 06, 1988

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DROPERIDOL

INJECTABLE; INJECTION

DROPERIDOL

SOLOPAK	2.5MG/ML	A071754	001	Sep 06, 1988
	2.5MG/ML	A071755	001	Sep 06, 1988
WATSON LABS	2.5MG/ML	A073520	001	Nov 27, 1991
	2.5MG/ML	A073521	001	Nov 27, 1991
	2.5MG/ML	A073523	001	Nov 27, 1991
INAPSINE				
+ RISING	2.5MG/ML	N016796	001	

DROPERIDOL; FENTANYL CITRATE

INJECTABLE; INJECTION

FENTANYL CITRATE AND DROPERIDOL

ASTRAZENECA	2.5MG/ML;EQ 0.05MG BASE/ML	A072026	001	Apr 13, 1989
	2.5MG/ML;EQ 0.05MG BASE/ML	A072027	001	Apr 13, 1989
	2.5MG/ML;EQ 0.05MG BASE/ML	A072028	001	Apr 13, 1989
HOSPIRA	2.5MG/ML;EQ 0.05MG BASE/ML	A071982	001	May 04, 1988
INNOVAR				
EPIC PHARMA LLC	2.5MG/ML;EQ 0.05MG BASE/ML	N016049	001	

DROSPIRENONE

TABLET, CHEWABLE; ORAL

DROSPIRENONE

+ EXELTIS USA INC	3.5MG	N216285	001	Jun 29, 2022
-------------------	-------	---------	-----	--------------

DROSPIRENONE; ETHINYL ESTRADIOL

TABLET; ORAL

DROSPIRENONE AND ETHINYL ESTRADIOL

BARR	3MG;0.02MG	A078515	001	Mar 30, 2009
JUBILANT CADISTA	3MG;0.02MG	A209423	001	Dec 22, 2017
KYRA				
SUN PHARM	3MG;0.02MG	A202318	001	Jul 23, 2019

TABLET; ORAL-28

DROSPIRENONE AND ETHINYL ESTRADIOL

APOTEX	3MG;0.03MG	A205876	001	Sep 21, 2016
BARR	3MG;0.03MG	A077527	001	May 09, 2008
JUBILANT CADISTA	3MG;0.03MG	A210017	001	Sep 10, 2018
KEMEYA				
SUN PHARM	3MG;0.03MG	A202138	001	Mar 13, 2019

DROXIDOPA

CAPSULE; ORAL

DROXIDOPA

CHARTWELL RX	100MG	A214217	001	May 05, 2022
	200MG	A214217	002	May 05, 2022
	300MG	A214217	003	May 05, 2022
HIKMA	100MG	A212835	001	Feb 18, 2021
	200MG	A212835	002	Feb 18, 2021
	300MG	A212835	003	Feb 18, 2021
TEVA PHARMS USA INC	100MG	A213162	001	Feb 18, 2021
	200MG	A213162	002	Feb 18, 2021
	300MG	A213162	003	Feb 18, 2021
UPSHER SMITH LABS	100MG	A213661	001	Feb 18, 2021
	200MG	A213661	002	Feb 18, 2021
	300MG	A213661	003	Feb 18, 2021

DULOXETINE HYDROCHLORIDE

CAPSULE, DELAYED REL PELLETS; ORAL

DULOXETINE HYDROCHLORIDE

APOTEX	EQ 20MG BASE	A202045	001	Jun 11, 2014
	EQ 30MG BASE	A202045	002	Jun 11, 2014
	EQ 60MG BASE	A202045	003	Jun 11, 2014
ENDO OPERATIONS	EQ 20MG BASE	A090780	001	Oct 28, 2015
	EQ 30MG BASE	A090780	002	Oct 28, 2015
	EQ 60MG BASE	A090780	003	Oct 28, 2015
MARKSANS PHARMA	EQ 20MG BASE	A090723	001	Dec 11, 2013
	EQ 30MG BASE	A090723	002	Dec 11, 2013
	EQ 60MG BASE	A090723	003	Dec 11, 2013
SUN PHARM	EQ 20MG BASE	A090745	001	Dec 11, 2013
	EQ 30MG BASE	A090745	002	Dec 11, 2013
	EQ 60MG BASE	A090745	003	Dec 11, 2013

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DULOXETINE HYDROCHLORIDE

CAPSULE, DELAYED REL PELLETS;ORAL

DULOXETINE HYDROCHLORIDE

TEVA PHARMS USA	EQ 20MG BASE	A090783 001	Dec 11, 2013
	EQ 30MG BASE	A090783 002	Dec 11, 2013
	EQ 60MG BASE	A090783 003	Dec 11, 2013
YAOPHARMA CO LTD	EQ 20MG BASE	A207219 001	Aug 16, 2019
	EQ 30MG BASE	A207219 002	Aug 16, 2019
	EQ 60MG BASE	A207219 003	Aug 16, 2019

DUTASTERIDE

CAPSULE;ORAL

DUTASTERIDE

ACTAVIS LABS FL INC	0.5MG	A202808 001	Nov 20, 2015
APOTEX	0.5MG	A204292 001	Nov 24, 2015
HERITAGE PHARMS INC	0.5MG	A207935 001	Oct 13, 2017
HIKMA	0.5MG	A202204 001	Nov 23, 2015
NORVIUM BIOSCIENCE	0.5MG	A203241 001	Jun 14, 2016
NOSTRUM LABS INC	0.5MG	A204705 001	Nov 20, 2015
RISING	0.5MG	A202530 001	Nov 20, 2015
STRIDES PHARMA	0.5MG	A208227 001	Jun 22, 2018
VINTAGE	0.5MG	A202421 001	Nov 20, 2015

DUTASTERIDE; TAMSULOSIN HYDROCHLORIDE

CAPSULE;ORAL

DUTASTERIDE AND TAMSULOSIN HYDROCHLORIDE

ACTAVIS LABS FL INC	0.5MG;0.4MG	A202975 001	Nov 20, 2015
---------------------	-------------	-------------	--------------

DYCLONINE HYDROCHLORIDE

SOLUTION;TOPICAL

DYCLONE

+ ASTRAZENECA	0.5% **	N009925 002	
+	1% **	N009925 001	

DYDROGESTERONE

TABLET;ORAL

GYNOREST

+ SOLVAY	5MG **	N017388 001	
+	10MG **	N017388 002	

DYPHYLLINE

ELIXIR;ORAL

NEOTHYLLINE

TEVA	160MG/15ML	N007794 003	
------	------------	-------------	--

INJECTABLE; INJECTION

NEOTHYLLINE

TEVA	250MG/ML	N009088 001	
------	----------	-------------	--

TABLET;ORAL

DILOR

SAVAGE LABS	200MG	A084514 001	
-------------	-------	-------------	--

DILOR-400

SAVAGE LABS	400MG	A084751 001	
-------------	-------	-------------	--

LUFYLLIN

NORVIUM BIOSCIENCE	200MG	A084566 001	
--------------------	-------	-------------	--

	400MG	A084566 002	
--	-------	-------------	--

NEOTHYLLINE

TEVA	200MG	N007794 001	
------	-------	-------------	--

	400MG	N007794 002	
--	-------	-------------	--

ECHOTHIOPHATE IODIDE

FOR SOLUTION;OPHTHALMIC

PHOSPHOLINE IODIDE

FERA PHARMS LLC	0.03%	N011963 002	
	0.06%	N011963 004	
	0.25%	N011963 003	

ECONAZOLE NITRATE

CREAM;TOPICAL

ECONAZOLE NITRATE

AUROBINDO PHARMA USA	1%	A210364 001	Apr 18, 2018
CHARTWELL RX	1%	A076075 001	Nov 26, 2002
ENCUBE	1%	A076574 001	Dec 17, 2004

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ECONAZOLE NITRATE

CREAM; TOPICAL

SPECTAZOLE

+ ALVOGEN

1% **

N018751 001 Dec 23, 1982

EDETATE CALCIUM DISODIUM

INJECTABLE; INJECTION

CALCIUM DISODIUM VERSENATE

+ BAUSCH

200MG/ML **

N008922 001

TABLET; ORAL

CALCIUM DISODIUM VERSENATE

BAUSCH

500MG

N008922 002

EDROPHONIUM CHLORIDE

INJECTABLE; INJECTION

EDROPHONIUM CHLORIDE

HOSPIRA

10MG/ML

A040131 001 Feb 24, 1998

WATSON LABS

10MG/ML

A040044 001 Mar 20, 1996

EDROPHONIUM CHLORIDE PRESERVATIVE FREE

WATSON LABS

10MG/ML

A040043 001 Mar 20, 1996

ENLON

NORVIUM BIOSCIENCE

10MG/ML

A088873 001 Aug 06, 1985

REVERSOL

ORGANON USA INC

10MG/ML

A089624 001 May 13, 1988

TENSILON

+ PAI HOLDINGS PHARM

10MG/ML **

N007959 001

TENSILON PRESERVATIVE FREE

+ PAI HOLDINGS PHARM

10MG/ML **

N007959 002

EFAVIRENZ

CAPSULE; ORAL

SUSTIVA

+ BRISTOL MYERS SQUIBB

50MG **

N020972 001 Sep 17, 1998

+ 100MG **

N020972 002 Sep 17, 1998

+ 200MG **

N020972 003 Sep 17, 1998

TABLET; ORAL

EFAVIRENZ

AUROBINDO PHARMA

600MG

A205322 001 Aug 30, 2018

MYLAN

600MG

A091471 001 Feb 17, 2016

STRIDES PHARMA

600MG

A078509 001 Jun 16, 2021

SUSTIVA

+ BRISTOL MYERS SQUIBB

300MG **

N021360 001 Feb 01, 2002

+ 600MG

N021360 002 Feb 01, 2002

EFAVIRENZ; EMTRICITABINE; TENOFOVIR DISOPROXIL FUMARATE

TABLET; ORAL

ATRIPLA

+ GILEAD SCIENCES

600MG; 200MG; 300MG

N021937 001 Jul 12, 2006

EFAVIRENZ; EMTRICITABINE AND TENOFOVIR DISOPROXIL FUMARATE

STRIDES PHARMA

600MG; 200MG; 300MG

A201802 001 Oct 03, 2023

EFAVIRENZ; EMTRICITABINE; TENOFOVIR DISOPROXIL FUMARATE

MYLAN

600MG; 200MG; 300MG

A209061 001 Sep 05, 2024

EFAVIRENZ; LAMIVUDINE; TENOFOVIR DISOPROXIL FUMARATE

TABLET; ORAL

EFAVIRENZ, LAMIVUDINE AND TENOFOVIR DISOPROXIL FUMARATE

+ AUROBINDO PHARMA LTD

600MG; 300MG; 300MG

N022343 001 Aug 15, 2018

MACLEODS PHARMS LTD

400MG; 300MG; 300MG

N210649 001 Mar 15, 2019

EFINACONAZOLE

SOLUTION; TOPICAL

EFINACONAZOLE

AUROBINDO PHARMA LTD

10%

A212066 001 Mar 29, 2021

LUPIN LTD

10%

A212169 001 Mar 02, 2022

PADAGIS US

10%

A211851 001 Dec 16, 2020

ZYDUS LIFESCIENCES

10%

A212178 001 Jul 15, 2022

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

EFLOARNITHINE HYDROCHLORIDE

CREAM; TOPICAL

VANIQA

+ SKINMEDICA

13.9%

N021145 001 Jul 27, 2000

INJECTABLE; INJECTION

ORNIDYL

SANOFI AVENTIS US

200MG/ML

N019879 002 Nov 28, 1990

ELETRIPTAN HYDROBROMIDE

TABLET; ORAL

ELETRIPTAN HYDROBROMIDE

STEVENS J

EQ 20MG BASE

A206787 001 May 25, 2018

EQ 40MG BASE

A206787 002 May 25, 2018

YUNG SHIN PHARM

EQ 20MG BASE

A209680 001 Jul 13, 2020

EQ 40MG BASE

A209680 002 Jul 13, 2020

ELIGLUSTAT TARTRATE

CAPSULE; ORAL

ELIGLUSTAT TARTRATE

DR REDDYS

EQ 84MG BASE

A212449 001 Aug 17, 2022

TEVA PHARMS USA INC

EQ 84MG BASE

A212474 001 Dec 27, 2021

ELTROMBOPAG OLAMINE

TABLET; ORAL

PROMACTA

+ NOVARTIS

EQ 100MG ACID **

N022291 005 Nov 16, 2012

ELVITEGRAVIR

TABLET; ORAL

VITEKTA

+ GILEAD SCIENCES INC

85MG

N203093 001 Sep 24, 2014

+

150MG

N203093 002 Sep 24, 2014

EMEDASTINE DIFUMARATE

SOLUTION/DROPS; OPHTHALMIC

EMADINE

+ NOVARTIS

0.05%

N020706 001 Dec 29, 1997

EMPAGLIFLOZIN

TABLET; ORAL

EMPAGLIFLOZIN

ZYDUS PHARMS

10MG

A212138 001 Aug 03, 2022

25MG

A212138 002 Aug 03, 2022

EMPAGLIFLOZIN; METFORMIN HYDROCHLORIDE

TABLET; ORAL

EMPAGLIFLOZIN AND METFORMIN HYDROCHLORIDE

ZYDUS PHARMS

5MG; 500MG

A212198 001 Jul 07, 2022

5MG; 1GM

A212198 002 Jul 07, 2022

12.5MG; 500MG

A212198 003 Jul 07, 2022

12.5MG; 1GM

A212198 004 Jul 07, 2022

EMTRICITABINE; TENOFOVIR ALAFENAMIDE FUMARATE

TABLET; ORAL

EMTRICITABINE AND TENOFOVIR ALAFENAMIDE FUMARATE

APOTEX

120MG; EQ 15MG BASE

A214053 001 May 17, 2024

200MG; EQ 25MG BASE

A214053 002 May 17, 2024

EMTRICITABINE; TENOFOVIR DISOPROXIL FUMARATE

TABLET; ORAL

EMTRICITABINE AND TENOFOVIR DISOPROXIL FUMARATE

APOTEX

200MG; 300MG

A208740 001 Jun 16, 2021

CIPLA

200MG; 300MG

A090958 001 Apr 02, 2021

ENALAPRIL MALEATE

FOR SOLUTION; ORAL

EPANED KIT

+ AZURITY

1MG/ML **

N204308 001 Aug 13, 2013

TABLET; ORAL

ENALAPRIL MALEATE

APOTHECON

2.5MG

A075583 001 Aug 22, 2000

5MG

A075583 002 Aug 22, 2000

10MG

A075583 003 Aug 22, 2000

20MG

A075583 004 Aug 22, 2000

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ENALAPRIL MALEATE

TABLET; ORAL

ENALAPRIL MALEATE

AUROBINDO PHARMA USA	2.5MG	A075480 001	Aug 22, 2000
	5MG	A075480 002	Aug 22, 2000
	10MG	A075480 003	Aug 22, 2000
	20MG	A075480 004	Aug 22, 2000
BEXIMCO PHARMS USA	2.5MG	A075621 001	Aug 22, 2000
	5MG	A075621 002	Aug 22, 2000
	10MG	A075621 003	Aug 22, 2000
	20MG	A075621 004	Aug 22, 2000
CHARTWELL RX	2.5MG	A075048 001	Aug 22, 2000
	5MG	A075048 002	Aug 22, 2000
	10MG	A075048 003	Aug 22, 2000
	20MG	A075048 004	Aug 22, 2000
IVAX SUB TEVA PHARMS	2.5MG	A075482 001	Aug 22, 2000
	5MG	A075482 002	Aug 22, 2000
	10MG	A075482 003	Aug 22, 2000
	20MG	A075482 004	Aug 22, 2000
KRKA DD NOVO MESTO	2.5MG	A075370 001	Aug 22, 2000
	5MG	A075370 002	Aug 22, 2000
	10MG	A075369 001	Aug 22, 2000
	20MG	A075369 002	Aug 22, 2000
MYLAN	2.5MG	A075472 001	Aug 22, 2000
	5MG	A075472 002	Aug 22, 2000
	10MG	A075472 003	Aug 22, 2000
	20MG	A075472 004	Aug 22, 2000
SUN PHARM INDS LTD	2.5MG	A075556 001	Aug 22, 2000
	5MG	A075556 002	Aug 22, 2000
	10MG	A075556 003	Aug 22, 2000
	20MG	A075556 004	Aug 22, 2000
WATSON LABS	2.5MG	A075501 001	Aug 22, 2000
	5MG	A075501 002	Aug 22, 2000
	10MG	A075501 003	Aug 22, 2000
	20MG	A075501 004	Aug 22, 2000

ENALAPRIL MALEATE; FELODIPINE

TABLET, EXTENDED RELEASE; ORAL

LEXXEL

ASTRAZENECA	5MG; 2.5MG	N020668 002	Oct 28, 1998
	5MG; 5MG	N020668 001	Dec 27, 1996

ENALAPRIL MALEATE; HYDROCHLOROTHIAZIDE

TABLET; ORAL

ENALAPRIL MALEATE AND HYDROCHLOROTHIAZIDE

CHARTWELL RX	5MG; 12.5MG	A076116 001	Sep 19, 2001
	10MG; 25MG	A076116 002	Sep 19, 2001
COSETTE	5MG; 12.5MG	A075727 001	Sep 18, 2001
	10MG; 25MG	A075727 002	Sep 18, 2001
IVAX SUB TEVA PHARMS	5MG; 12.5MG	A075736 001	Mar 25, 2003
	10MG; 25MG	A075736 002	Mar 25, 2003
NOSTRUM LABS INC	5MG; 12.5MG	A076486 001	Oct 27, 2004
	10MG; 25MG	A076486 002	Oct 27, 2004
RISING	5MG; 12.5MG	A075624 001	Sep 18, 2001
	10MG; 25MG	A075624 002	Sep 18, 2001

ENALAPRILAT

INJECTABLE; INJECTION

ENALAPRILAT

HOSPIRA	1.25MG/ML	A075456 001	Aug 22, 2000
	1.25MG/ML	A075571 001	Aug 22, 2000
VASOTEC			
+ BIOVAIL LABS INTL	1.25MG/ML **	N019309 001	Feb 09, 1988

ENCORAFENIB

CAPSULE; ORAL

BRAFTOVI

+ ARRAY BIOPHARMA INC	50MG	N210496 001	Jun 27, 2018
-----------------------	------	-------------	--------------

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ENFLURANE

LIQUID; INHALATION

ENFLURANE

ABBOTT	99.9%	A070803	001	Sep 08, 1987
PIRAMAL CRITICAL	99.9%	A074396	001	Jul 29, 1994
ETHRANE				
BAXTER HLTHCARE	99.9% **	N017087	001	

ENOXACIN

TABLET; ORAL

PENETREX

SANOFI AVENTIS US	200MG	N019616	004	Dec 31, 1991
	400MG	N019616	005	Dec 31, 1991

ENOXAPARIN SODIUM

INJECTABLE; SUBCUTANEOUS

LOVENOX (PRESERVATIVE FREE)

+ SANOFI AVENTIS US	90MG/0.6ML (150MG/ML) **	N020164	006	Jun 02, 2000
---------------------	--------------------------	---------	-----	--------------

ENTACAPONE

TABLET; ORAL

ENTACAPONE

NORVIUM BIOSCIENCE	200MG	A202394	001	May 13, 2013
--------------------	-------	---------	-----	--------------

ENTECAVIR

TABLET; ORAL

ENTECAVIR

ACCORD HLTHCARE	0.5MG	A205824	001	Aug 25, 2017
	1MG	A205824	002	Aug 25, 2017
CHARTWELL RX	0.5MG	A206294	001	Nov 23, 2016
	0.5MG	A206672	001	May 11, 2017
	1MG	A206294	002	Nov 23, 2016
	1MG	A206672	002	May 11, 2017
PRINSTON INC	0.5MG	A208782	001	Oct 10, 2017
	1MG	A208782	002	Oct 10, 2017
RISING	0.5MG	A206226	001	Mar 26, 2019
	1MG	A206226	002	Mar 26, 2019
SUNSHINE	0.5MG	A211978	001	May 20, 2020
	1MG	A211978	002	May 20, 2020
SWISS PHARM	0.5MG	A212106	001	Aug 10, 2020
	1MG	A212106	002	Aug 10, 2020
TEVA PHARMS USA	0.5MG	A202122	001	Aug 26, 2014
	1MG	A202122	002	Aug 26, 2014
YAOPHARMA CO LTD	0.5MG	A212201	001	Nov 04, 2019
	1MG	A212201	002	Nov 04, 2019

ENZALUTAMIDE

CAPSULE; ORAL

ENZALUTAMIDE

ACTAVIS LABS FL INC	40MG	A209614	001	May 14, 2021
APOTEX	40MG	A209645	001	Apr 22, 2022

EPHEDRINE HYDROCHLORIDE

SOLUTION; INTRAVENOUS

REZIPRES

+ DR REDDYS LABS SA	23.5MG/5ML (4.7MG/ML)	N213536	001	Jun 14, 2021
+	47MG/ML (47MG/ML)	N213536	002	Jun 14, 2021
+	47MG/5ML (9.4MG/ML)	N213536	003	Jun 14, 2021

EPHEDRINE SULFATE

SOLUTION; INTRAVENOUS

EPHEDRINE SULFATE

RENEW PHARMS	50MG/ML (50MG/ML)	N208609	001	Mar 01, 2017
ZYDUS PHARMS	50MG/ML (50MG/ML)	A217276	001	May 16, 2023

EPINASTINE HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

ELESTAT

+ ALLERGAN	0.05% **	N021565	001	Oct 16, 2003
EPINASTINE HYDROCHLORIDE				
BRECKENRIDGE	0.05%	A090870	001	Mar 14, 2011
CHARTWELL RX	0.05%	A203384	001	Dec 07, 2016
EPIC PHARMA LLC	0.05%	A204055	001	May 05, 2017

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

EPINASTINE HYDROCHLORIDESOLUTION/DROPS;OPHTHALMIC
EPINASTINE HYDROCHLORIDE
SUN PHARM INDS

0.05%

A091626 001 Oct 31, 2011

EPINEPHRINEAEROSOL, METERED; INHALATION
BRONKAID MIST

STERLING

0.25MG/INH

N016803 001

EPINEPHRINE

ARMSTRONG PHARMS

0.2MG/INH

A087907 001 May 23, 1984

PRIMATENE MIST

WYETH CONS

0.2MG/INH

N016126 001

INJECTABLE; INJECTION

SUS-PHRINE SULFITE FREE

FOREST LABS

1.5MG/AMP
5MG/MLN007942 003 Feb 05, 1999
N007942 001

INJECTABLE; INTRAMUSCULAR

EPI E Z PEN JR

MYLAN SPECIALITY LP

0.15MG/DELIVERY

N019430 004 Aug 03, 1995

EPIPEN E Z PEN

MYLAN SPECIALITY LP

0.3MG/DELIVERY

N019430 003 Aug 03, 1995

INJECTABLE; INTRAMUSCULAR, SUBCUTANEOUS

TWINJECT 0.15

IMPAX

EQ 0.15MG/DELIVERY

N020800 002 May 28, 2004

TWINJECT 0.3

IMPAX

EQ 0.3MG/DELIVERY

N020800 001 May 30, 2003

SOLUTION; INTRAMUSCULAR, INTRAVENOUS, SUBCUTANEOUS

EPINEPHRINE

AM REGENT

EQ 1MG BASE/ML (EQ 1MG BASE/ML)

A207568 001 Jul 06, 2018

SOLUTION; INTRAMUSCULAR, SUBCUTANEOUS

SYMJEPI

+ ADAMIS PHARMS CORP

0.15MG/0.3ML (0.15MG/0.3ML)

N207534 002 Sep 27, 2018

SOLUTION; INTRAVENOUS

EPINEPHRINE

+ BPI LABS

1MG/ML (1MG/ML)

N205029 003 May 12, 2023

EPINEPHRINE BITARTRATEAEROSOL, METERED; INHALATION
BRONITIN MIST

WYETH CONS

0.3MG/INH

N016126 002

MEDIHALER-EPI

3M

0.3MG/INH

N010374 003

EPINEPHRINE BITARTRATE; ETIDOCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

DURANEST

+ ASTRAZENECA

0.005MG/ML; 1% **

N017751 006

+

0.005MG/ML; 1.5% **

N017751 007

+ DENTSPLY PHARM

0.005MG/ML; 1.5% **

N021384 001

EPINEPHRINE BITARTRATE; PRILOCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

CITANEST FORTE

ASTRAZENECA

0.005MG/ML; 4%

N014763 008

EPINEPHRINE; ETIDOCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

DURANEST

+ ASTRAZENECA

0.005MG/ML; 0.5% **

N017751 004

EPINEPHRINE; LIDOCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

ALPHACAINE HYDROCHLORIDE W/ EPINEPHRINE

CARLISLE

0.01MG/ML; 2%

A084720 001

0.02MG/ML; 2%

A084732 001

LIDOCAINE HYDROCHLORIDE AND EPINEPHRINE

BELMORA LLC

0.01MG/ML; 2%

A080504 004 Oct 19, 1983

0.02MG/ML; 2%

A080504 005 Oct 19, 1983

EASTMAN KODAK

0.01MG/ML; 2%

A040057 002 Feb 26, 1993

0.02MG/ML; 2%

A040057 001 Feb 26, 1993

HOSPIRA

0.005MG/ML; 1%

A089649 001 Jun 21, 1988

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

EPINEPHRINE; LIDOCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

LIDOCAINE HYDROCHLORIDE AND EPINEPHRINE				
	0.005MG/ML;1.5%	A089650	001	Jun 21, 1988
	0.01MG/ML;2%	A078772	001	May 12, 2008
	0.02MG/ML;2%	A078772	002	May 12, 2008
WEST-WARD PHARMS INT	0.01MG/ML;1%	A080406	001	
	0.01MG/ML;2%	A080406	002	
LIDOCAINE HYDROCHLORIDE W/ EPINEPHRINE				
ABBOT	0.01MG/ML;1%	A083154	001	
BEL MAR	0.01MG/ML;1%	A080820	001	
	0.01MG/ML;2%	A080757	001	
DELL LABS	0.01MG/ML;1%	A083389	001	
	0.01MG/ML;2%	A083390	001	
INTL MEDICATION	0.01MG/ML;1%	A086402	001	
WATSON LABS	0.01MG/ML;1%	A080377	003	
	0.01MG/ML;1%	A085463	001	
	0.01MG/ML;2%	A080377	004	
LIDOCATON				
PHARMATON	0.01MG/ML;2%	A084729	001	Aug 17, 1983
	0.02MG/ML;2%	A084728	001	Aug 17, 1983
OCTOCAINE				
SEPTODONT	0.01MG/ML;2%	A084048	001	
	0.02MG/ML;2%	A084048	002	
XYLOCAINE DENTAL WITH EPINEPHRINE				
DENTSPLY PHARM	0.01MG/ML;2% **	N021381	001	
	0.02MG/ML;2% **	N021381	002	
XYLOCAINE W/ EPINEPHRINE				
ASTRAZENECA	0.005MG/ML;1%	N010418	006	
	0.005MG/ML;1.5%	N010418	010	
	0.005MG/ML;2%	N010418	008	
+ FRESENIUS KABI USA	0.01MG/ML;2%	N006488	003	
PATCH; IONTOPHORESIS, TOPICAL				
LIDOSITE TOPICAL SYSTEM KIT				
VYTERIS	1.05MG/PATCH;100MG/PATCH	N021504	001	May 06, 2004
SOLUTION; IONTOPHORESIS				
IONTOCAINE				
IOMED	0.01MG/ML;2%	N020530	001	Dec 21, 1995
SOLUTION; IONTOPHORESIS, TOPICAL				
LIDOCAINE HYDROCHLORIDE AND EPINEPHRINE				
EMPI	0.01MG/ML;2%	N021486	001	Oct 26, 2004

EPINEPHRINE; PROCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

PROCAINE HYDROCHLORIDE W/ EPINEPHRINE				
BEL MAR	0.02MG/ML;1%	A080758	001	
	0.02MG/ML;2%	A080759	001	

EPIRUBICIN HYDROCHLORIDE

INJECTABLE; INJECTION

EPIRUBICIN HYDROCHLORIDE				
ACTAVIS TOTOWA	10MG/5ML (2MG/ML)	A065445	001	Sep 18, 2008
	50MG/25ML (2MG/ML)	A065445	002	Sep 18, 2008
	200MG/100ML (2MG/ML)	A065445	003	Sep 18, 2008
EBEWE PHARMA	50MG/25ML (2MG/ML)	A065339	001	Dec 22, 2009
	200MG/100ML (2MG/ML)	A065339	002	Dec 22, 2009
FRESENIUS KABI USA	10MG/5ML (2MG/ML)	A065408	001	Oct 15, 2007
	50MG/25ML (2MG/ML)	A065408	002	Oct 15, 2007
	150MG/75ML (2MG/ML)	A065408	003	Oct 15, 2007
	200MG/100ML (2MG/ML)	A065408	004	Oct 15, 2007
	200MG/100ML (2MG/ML)	A065411	001	Aug 20, 2007
	50MG/25ML (2MG/ML)	A065411	002	Aug 20, 2007
HOSPIRA	10MG/5ML (2MG/ML)	A065343	001	Apr 19, 2007
	50MG/25ML (2MG/ML)	A065343	002	Apr 19, 2007
	150MG/75ML (2MG/ML)	A065343	003	Apr 19, 2007
	200MG/100ML (2MG/ML)	A065343	004	Apr 19, 2007
NORVIUM BIOSCIENCE	50MG/25ML (2MG/ML)	A065371	001	Nov 28, 2007
	50MG/25ML (2MG/ML)	A091599	001	Mar 12, 2012
	200MG/100ML (2MG/ML)	A065371	002	Nov 28, 2007
	200MG/100ML (2MG/ML)	A091599	002	Mar 12, 2012

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

EPIRUBICIN HYDROCHLORIDE

INJECTABLE; INJECTION

EPIRUBICIN HYDROCHLORIDE

ZENNOVA

50MG/25ML (2MG/ML)

A090266 001 Apr 15, 2011

200MG/100ML (2MG/ML)

A090266 002 Apr 15, 2011

POWDER; INTRAVENOUS

EPIRUBICIN HYDROCHLORIDE

HOSPIRA

50MG/VIAL

N050807 001 Sep 15, 2006

200MG/VIAL

N050807 002 Sep 15, 2006

EPLERENONE

TABLET; ORAL

EPLERENONE

PRASCO

25MG

A203896 001 Feb 02, 2017

50MG

A203896 002 Feb 02, 2017

INSPRA

UPJOHN

100MG

N021437 003 Sep 27, 2002

EPROSARTAN MESYLATE

TABLET; ORAL

EPROSARTAN MESYLATE

NORVIUM BIOSCIENCE

EQ 400MG BASE

A202012 001 Nov 16, 2011

EQ 600MG BASE

A202012 002 Nov 16, 2011

TEVETEN

ABBVIE

EQ 300MG BASE **

N020738 004 Dec 22, 1997

+

EQ 400MG BASE **

N020738 005 Dec 22, 1997

+

EQ 600MG BASE **

N020738 006 May 27, 1999

EPROSARTAN MESYLATE; HYDROCHLOROTHIAZIDE

TABLET; ORAL

TEVETEN HCT

ABBVIE

600MG; 12.5MG

N021268 001 Nov 01, 2001

600MG; 25MG

N021268 002 Nov 01, 2001

EPTIFIBATIDE

INJECTABLE; INJECTION

EPTIFIBATIDE

ACCORD HLTHCARE

2MG/ML

A205557 001 Nov 06, 2017

75MG/100ML

A205557 002 Nov 06, 2017

AMNEAL PHARMS

2MG/ML

A205581 001 Dec 08, 2016

75MG/100ML

A205581 002 Dec 08, 2016

BAXTER HLTHCARE CORP

2MG/ML

A208554 001 Nov 23, 2018

HYBIO

2MG/ML

A207864 001 Mar 20, 2020

75MG/100ML

A207864 002 Mar 20, 2020

RISING

2MG/ML

A204589 001 Apr 18, 2017

75MG/100ML

A204589 002 Apr 18, 2017

TEVA PHARMS USA

75MG/100ML

A091555 001 Jun 05, 2015

USV

2MG/ML

A204361 001 Mar 14, 2019

2MG/ML

A204362 001 Mar 11, 2019

75MG/100ML

A204361 002 Mar 14, 2019

INTEGRILIN

+

MSD SUB MERCK

2MG/ML **

N020718 001 May 18, 1998

+

75MG/100ML **

N020718 002 May 18, 1998

ERGOCALCIFEROL

CAPSULE; ORAL

DELTALIN

LILLY

50,000 IU

A080884 001

ERGOCALCIFEROL

SIGMAPHARM LABS LLC

50,000 IU

A091004 001 Jul 14, 2010

SUN PHARM INDS INC

50,000 IU

A040865 001 Dec 29, 2009

VITAMIN D

CHARTWELL MOLECULAR

50,000 IU

A080825 001

CHASE CHEM

50,000 IU

A080747 001

EVERYLIFE

50,000 IU

A080956 001

IMPAX LABS

50,000 IU

A080951 001

VITARINE

50,000 IU

A084053 001

WEST WARD

50,000 IU

A083102 001

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ERGOLOID MESYLATES

CAPSULE; ORAL

HYDERGINE LC

NOVARTIS

1MG

N018706 001 Jan 18, 1983

SOLUTION; ORAL

HYDERGINE

NOVARTIS

1MG/ML

N018418 001

TABLET; ORAL

ERGOLOID MESYLATES

MUTUAL PHARM

1MG

A088891 001 Nov 01, 1985

SUN PHARM INDUSTRIES

1MG

A081113 001 Oct 31, 1991

WATSON LABS

1MG

A086433 001 May 27, 1982

1MG

A087244 001 Aug 16, 1982

GERIMAL

WATSON LABS

1MG

A088207 001 Mar 22, 1984

HYDERGINE

NOVARTIS

0.5MG **

N017993 003

+

1MG **

N017993 001

TABLET; SUBLINGUAL

ALKERGOT

SANDOZ

0.5MG

A085153 001

1MG

A087417 001

CIRCANOL

3M

0.5MG

A084868 001

1MG

A085809 001

DEAPRIL-ST

BRISTOL MYERS SQUIBB

1MG

A085020 002

ERGOLOID MESYLATES

KV PHARM

0.5MG

A085899 001

0.5MG

A086265 001

1MG

A085900 001

1MG

A086264 001

LEDERLE

0.5MG

A086984 001

1MG

A086985 001

SUN PHARM INDUSTRIES

0.5MG

A087407 001

1MG

A087552 001

SUPERPHARM

0.5MG

A089233 001 Sep 23, 1986

1MG

A089234 001 Sep 23, 1986

VANGARD

0.5MG

A088013 001 Sep 20, 1982

1MG

A088014 001 Sep 20, 1982

WATSON LABS

0.5MG

A084930 001

0.5MG

A087233 001

1MG

A085177 001

1MG

A087183 001

GERIMAL

WATSON LABS

0.5MG

A086189 001

1MG

A086188 001

HYDERGINE

NOVARTIS

0.5MG

N009087 002

1MG

N009087 001

HYDROGENATED ERGOT ALKALOIDS

IVAX PHARMS

0.5MG

A087186 001

1MG

A087185 001

ERGOTAMINE TARTRATE

AEROSOL, METERED; INHALATION

MEDIHALER ERGOTAMINE

3M

0.36MG/INH

N012102 001

TABLET; SUBLINGUAL

ERGOSTAT

WATSON LABS INC

2MG

A088337 001 Jun 08, 1984

WIGRETTES

ORGANON USA INC

2MG

A086750 001 Jul 29, 1982

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ERLOTINIB HYDROCHLORIDE

TABLET;ORAL

ERLOTINIB HYDROCHLORIDE

ACCORD HLTHCARE	EQ 25MG BASE	A211083 001	Jul 02, 2020
	EQ 100MG BASE	A211083 002	Jul 02, 2020
	EQ 150MG BASE	A211083 003	Jul 02, 2020
EUGIA PHARMA	EQ 25MG BASE	A216342 001	Jun 22, 2022
	EQ 100MG BASE	A216342 002	Jun 22, 2022
	EQ 150MG BASE	A216342 003	Jun 22, 2022
NATCO PHARMA LTD	EQ 25MG BASE	A208488 001	Nov 05, 2019
	EQ 150MG BASE	A208488 003	Nov 05, 2019
TARCEVA			
+ OSI PHARMS	EQ 25MG BASE **	N021743 001	Nov 18, 2004
+	EQ 100MG BASE **	N021743 002	Nov 18, 2004
+	EQ 150MG BASE **	N021743 003	Nov 18, 2004

ERTAPENEM SODIUM

INJECTABLE; INTRAMUSCULAR, INTRAVENOUS

ERTAPENEM SODIUM

SUN PHARM	EQ 1GM BASE/VIAL	A209145 001	May 02, 2023
-----------	------------------	-------------	--------------

ERTUGLIFLOZIN

TABLET;ORAL

ERTUGLIFLOZIN

AUROBINDO PHARMA LTD	5MG	A216947 001	Jul 13, 2023
	15MG	A216947 002	Jul 13, 2023

ERYTHROMYCIN

CAPSULE, DELAYED REL PELLETS;ORAL

ERYC

PARKE DAVIS	250MG	A062546 001	Jul 25, 1985
	250MG	A062618 001	Sep 25, 1985
WARNER CHILCOTT LLC	250MG	A062338 001	

ERYC 125

PARKE DAVIS	125MG	A062648 001	Oct 24, 1985
-------------	-------	-------------	--------------

ERYC SPRINKLES

HOSPIRA	125MG	N050593 001	Jul 22, 1985
---------	-------	-------------	--------------

ERYTHROMYCIN

BARR	250MG	A063098 001	May 04, 1989
------	-------	-------------	--------------

GEL;TOPICAL

E-GLADES

NORVIUM BIOSCIENCE	2%	A065009 001	Mar 18, 2002
--------------------	----	-------------	--------------

EMGEL

ALTANA	2%	A063107 001	Aug 23, 1991
--------	----	-------------	--------------

ERYTHROMYCIN

ENCUBE	2%	A208154 001	Jul 19, 2017
--------	----	-------------	--------------

LOTION;TOPICAL

E-SOLVE 2

SYOSSET	2%	A062467 001	Jul 03, 1985
---------	----	-------------	--------------

OINTMENT;OPHTHALMIC

ERYTHROMYCIN

PADAGIS US	0.5%	A062447 001	Sep 26, 1983
------------	------	-------------	--------------

PHARMADERM	5MG/GM	A062446 001	Sep 26, 1983
------------	--------	-------------	--------------

PHARMAFAIR	5MG/GM	A062481 001	Apr 05, 1984
------------	--------	-------------	--------------

ILOTYCIN

DISTA	0.5% **	N050368 001	
-------	---------	-------------	--

OINTMENT;TOPICAL

AKNE-MYCIN

+ BAUSCH	2%	N050584 001	Jan 10, 1985
----------	----	-------------	--------------

POWDER;FOR RX COMPOUNDING

ERYTHROMYCIN

PADDOCK LLC	100%	N050610 001	Nov 07, 1986
-------------	------	-------------	--------------

SOLUTION;TOPICAL

A/T/S

TARO	2%	A062405 001	Nov 18, 1982
------	----	-------------	--------------

C-SOLVE-2

FOUGERA PHARMS	2%	A062468 001	Jul 03, 1985
----------------	----	-------------	--------------

ERYDERM

ARBOR PHARMS INC	2%	A062290 001	
------------------	----	-------------	--

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ERYTHROMYCIN

SOLUTION; TOPICAL

ERYMAX					
MERZ PHARMS	2%		A062508	002	Jul 11, 1985
ERYTHRO-STATIN					
EPIC PHARMA LLC	2%		A064101	001	Oct 22, 1996
ERYTHROMYCIN					
ALPHARMA US PHARMS	1.5%		A062328	001	Apr 19, 1982
	2%		A062326	001	Apr 19, 1982
	2%		A062327	001	Apr 19, 1982
	2%		A062342	001	Feb 25, 1982
	2%		A062957	001	Jul 21, 1988
BAUSCH	2%		A064039	001	Jan 27, 1994
FOUGERA PHARMS	2%		A064187	001	Sep 30, 1997
LILLY	2%		N050532	001	
PAI HOLDINGS PHARM	2%		A208100	001	Nov 20, 2017
PHARMAFAIR	1.5%		A062485	001	Jul 11, 1984
	2%		A062616	001	Jul 25, 1985
PHARMOBEDIANT CNSLTG	2%		A062825	001	Oct 23, 1987
RENAISSANCE PHARMA	2%		A064127	001	Feb 14, 1997
SANSAC					
DOW PHARM	2%		A062522	001	Jan 24, 1985
STATICIN					
+ WESTWOOD SQUIBB	1.5% **		N050526	001	
T-STAT					
WESTWOOD SQUIBB	2% **		A062436	001	Mar 09, 1983
SWAB; TOPICAL					
C-SOLVE-2					
IVAX SUB TEVA PHARMS	2%		A062751	001	Jul 30, 1993
ERYCETTE					
+ JOHNSON AND JOHNSON	2% **		N050594	001	Feb 15, 1985
ERYTHROMYCIN					
FOUGERA PHARMS	2%		A065320	001	Jul 25, 2006
NORVIUM BIOSCIENCE	2%		A064128	001	Jul 03, 1996
T-STAT					
WESTWOOD SQUIBB	2%		A062748	001	Jul 23, 1987
TABLET; ORAL					
ERYTHROMYCIN					
AMNEAL PHARMS CO	250MG		A209720	001	Mar 09, 2018
	500MG		A209720	002	Mar 09, 2018
ZYDUS PHARMS	250MG		A215440	001	Aug 31, 2023
	500MG		A215440	002	Aug 31, 2023
TABLET, COATED PARTICLES; ORAL					
PCE					
+ AZURITY	333MG		N050611	001	Sep 09, 1986
+	500MG		N050611	002	Aug 22, 1990
TABLET, DELAYED RELEASE; ORAL					
E-BASE					
BARR	333MG		A063028	001	May 15, 1990
	333MG		A063086	001	May 15, 1990
	500MG		A062999	001	Nov 25, 1988
E-MYCIN					
ARBOR PHARMS INC	250MG		A060272	001	
	333MG		A060272	002	
ILOTYCIN					
DISTA	250MG		A061910	001	
R-P MYCIN					
SOLVAY	250MG		A061659	001	
ROBIMYCIN					
ROBINS AH	250MG		A061633	001	

ERYTHROMYCIN ESTOLATE

CAPSULE; ORAL

ERYTHROMYCIN ESTOLATE					
BARR	EQ 125MG BASE		A062162	001	
	EQ 250MG BASE		A062162	002	
IVAX SUB TEVA PHARMS	EQ 250MG BASE		A062237	001	
WATSON LABS	EQ 250MG BASE		A062087	001	

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ERYTHROMYCIN ESTOLATE

CAPSULE; ORAL

ILOSONE

LILLY	EQ 125MG BASE	A061897 001	
	EQ 250MG BASE	A061897 002	

FOR SUSPENSION; ORAL

ILOSONE

DISTA	EQ 125MG BASE/5ML	A061893 001	
-------	-------------------	-------------	--

SUSPENSION; ORAL

ERYTHROMYCIN ESTOLATE

ALPHARMA US PHARMS	EQ 125MG BASE/5ML	A062353 001	Nov 18, 1982
	EQ 250MG BASE/5ML	A062409 001	Dec 16, 1982
COSETTE	EQ 125MG BASE/5ML	A062169 001	Oct 17, 1990
	EQ 250MG BASE/5ML	A062169 002	Oct 17, 1990
LIFE LABS	EQ 250MG BASE/5ML	A062362 001	Dec 17, 1982

ILOSONE

LILLY	EQ 125MG BASE/5ML	A061894 001	
	EQ 125MG BASE/5ML	N050010 001	
	EQ 250MG BASE/5ML	A061894 002	
	EQ 250MG BASE/5ML	N050010 002	

SUSPENSION/DROPS; ORAL

ILOSONE

LILLY	EQ 100MG BASE/ML	A061894 003	
-------	------------------	-------------	--

TABLET; ORAL

ILOSONE

LILLY	EQ 500MG BASE	A061896 001	
-------	---------------	-------------	--

TABLET, CHEWABLE; ORAL

ILOSONE

DISTA	EQ 125MG BASE	A061895 001	
	EQ 250MG BASE	A061895 002	

ERYTHROMYCIN ESTOLATE; SULFISOXAZOLE ACETYL

SUSPENSION; ORAL

ILOSONE SULFA

LILLY	EQ 125MG BASE/5ML; EQ 600MG BASE/5ML	N050599 001	Sep 29, 1989
-------	--------------------------------------	-------------	--------------

ERYTHROMYCIN ETHYLSUCCINATE

GRANULE; ORAL

ERYTHROMYCIN ETHYLSUCCINATE

ENDO OPERATIONS	EQ 200MG BASE/5ML	A211991 001	Oct 23, 2019
	EQ 400MG BASE/5ML	A211991 002	Oct 23, 2019

PEDIAMYCIN

ROSS LABS	EQ 200MG BASE/5ML	A062305 001	
-----------	-------------------	-------------	--

SUSPENSION; ORAL

E-MYCIN E

PHARMACIA AND UPJOHN	EQ 200MG BASE/5ML	A062198 001	
	EQ 400MG BASE/5ML	A062198 002	

E.E.S. 200

ARBOR PHARMS LLC	EQ 200MG BASE/5ML **	A061639 001	
------------------	----------------------	-------------	--

E.E.S. 400

ARBOR PHARMS LLC	EQ 400MG BASE/5ML **	A061639 002	
------------------	----------------------	-------------	--

ERYTHROMYCIN ETHYLSUCCINATE

ALPHARMA US PHARMS	EQ 200MG BASE/5ML	A062200 001	
	EQ 400MG BASE/5ML	A062200 002	
DISTA	EQ 200MG BASE/5ML	A062177 001	
	EQ 400MG BASE/5ML	A062177 002	
NASKA	EQ 400MG BASE/5ML	A062674 001	Mar 10, 1987
PARKE DAVIS	EQ 200MG BASE/5ML	A062231 001	
	EQ 400MG BASE/5ML	A062231 002	
PHARMAFAIR	EQ 200MG BASE/5ML	A062559 001	Mar 15, 1985
	EQ 400MG BASE/5ML	A062558 001	Mar 15, 1985

PEDIAMYCIN

ARBOR PHARMS LLC	EQ 200MG BASE/5ML	A062304 001	
------------------	-------------------	-------------	--

PEDIAMYCIN 400

ARBOR PHARMS LLC	EQ 400MG BASE/5ML	A062304 002	
------------------	-------------------	-------------	--

WYAMYCIN E

WYETH AYERST	EQ 200MG BASE/5ML	A062123 002	
	EQ 400MG BASE/5ML	A062123 001	

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ERYTHROMYCIN ETHYLSUCCINATE

SUSPENSION/DROPS;ORAL

PEDIAMYCIN

ROSS LABS EQ 100MG BASE/2.5ML A062305 002

TABLET;ORAL

E.E.S. 400

CARNEGIE EQ 400MG BASE A061905 001

ERYTHROMYCIN ETHYLSUCCINATE

AUROBINDO PHARMA USA EQ 400MG BASE A062847 001 Sep 14, 1988

BARR EQ 400MG BASE A062256 001

TABLET, CHEWABLE;ORAL

E.E.S.

AZURITY EQ 200MG BASE N050297 002

ERYPED

AZURITY EQ 200MG BASE N050297 003 Jul 05, 1988

PEDIAMYCIN

ROSS LABS EQ 200MG BASE A062306 001

ERYTHROMYCIN ETHYLSUCCINATE; SULFISOXAZOLE ACETYL

GRANULE;ORAL

ERYTHROMYCIN ETHYLSUCCINATE AND SULFISOXAZOLE ACETYL

BARR EQ 200MG BASE/5ML;EQ 600MG BASE/5ML A062759 001 May 20, 1988

ERYZOLE

ALRA EQ 200MG BASE/5ML;EQ 600MG BASE/5ML A062758 001 Jun 15, 1988

PEDIAZOLE

ROSS LABS EQ 200MG BASE/5ML;EQ 600MG BASE/5ML ** N050529 001

ERYTHROMYCIN GLUCEPTATE

INJECTABLE; INJECTION

ILOTYCIN GLUCEPTATE

DISTA EQ 250MG BASE/VIAL N050370 001

EQ 500MG BASE/VIAL N050370 002

EQ 1GM BASE/VIAL N050370 003

ERYTHROMYCIN LACTOBIONATE

INJECTABLE; INJECTION

ERYTHROCIN

ABBOTT EQ 500MG BASE/VIAL A062586 001 Jan 04, 1988

EQ 1GM BASE/VIAL A062586 002 Jan 04, 1988

HOSPIRA EQ 500MG BASE/VIAL N050182 002

EQ 1GM BASE/VIAL A062638 002 Oct 31, 1986

EQ 1GM BASE/VIAL N050182 003

+ EQ 1GM BASE/VIAL N050609 002 Sep 24, 1986

ERYTHROMYCIN

ELKINS SINN EQ 500MG BASE/VIAL A062563 001 Mar 28, 1985

EQ 1GM BASE/VIAL A062563 002 Mar 28, 1985

ERYTHROMYCIN LACTOBIONATE

ABRAXIS PHARM EQ 500MG BASE/VIAL A062604 001 Nov 24, 1986

EQ 1GM BASE/VIAL A062604 002 Nov 24, 1986

BAXTER HLTHCARE EQ 500MG BASE/VIAL A062993 001 May 09, 1989

EQ 1GM BASE/VIAL A062993 002 May 09, 1989

EXELA PHARMA EQ 500MG BASE/VIAL A211086 001 Sep 17, 2020

TEVA PARENTERAL EQ 500MG BASE/VIAL A063253 001 Jul 30, 1993

EQ 1GM BASE/VIAL A063253 002 Jul 30, 1993

ERYTHROMYCIN STEARATE

TABLET;ORAL

BRISTAMYCIN

BRISTOL EQ 250MG BASE A061304 001

EQ 250MG BASE A061887 001

ERYPAR

PARKE DAVIS EQ 250MG BASE A062032 001

EQ 500MG BASE A062032 002

WARNER CHILCOTT EQ 250MG BASE A062322 001

ERYTHROCIN STEARATE

AZURITY EQ 125MG BASE A060359 002

EQ 500MG BASE A060359 003

ERYTHROMYCIN STEARATE

ANI PHARMS EQ 250MG BASE A061461 001

EQ 250MG BASE A061591 001

EQ 500MG BASE A061461 002

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ERYTHROMYCIN STEARATE

TABLET; ORAL

ERYTHROMYCIN STEARATE

	EQ 500MG BASE	A063179 001	May 15, 1990
LEDERLE	EQ 250MG BASE	A062089 001	
	EQ 500MG BASE	A062089 002	
NORVIUM BIOSCIENCE	EQ 250MG BASE	A061505 001	
	EQ 500MG BASE	A061505 002	
PUREPAC PHARM	EQ 250MG BASE	A061743 001	
WATSON LABS	EQ 250MG BASE	A062121 002	
	EQ 500MG BASE	A062121 001	
ETHRIL 250			
BRISTOL MYERS SQUIBB	EQ 250MG BASE	A061605 001	
ETHRIL 500			
BRISTOL MYERS SQUIBB	EQ 500MG BASE	A061605 002	
PFIZER-E			
PFIZER	EQ 250MG BASE	A061791 001	
	EQ 500MG BASE	A061791 002	
WYAMYCIN S			
WYETH AYERST	EQ 250MG BASE	A061675 001	
	EQ 500MG BASE	A061675 002	

ESCITALOPRAM OXALATE

CAPSULE; ORAL

ESCITALOPRAM OXALATE

NORVIUM BIOSCIENCE	EQ 5MG BASE	A077660 001	Jul 31, 2007
	EQ 10MG BASE	A077660 002	Jul 31, 2007
	EQ 20MG BASE	A077660 003	Jul 31, 2007

SOLUTION; ORAL

ESCITALOPRAM OXALATE

ANTRIM PHARMS LLC	EQ 5MG BASE/5ML	A203967 001	May 26, 2015
-------------------	-----------------	-------------	--------------

LEXAPRO

+ ABBVIE

	EQ 5MG BASE/5ML **	N021365 001	Nov 27, 2002
--	--------------------	-------------	--------------

TABLET; ORAL

ESCITALOPRAM OXALATE

AIPING PHARM INC	EQ 5MG BASE	A077512 001	Sep 12, 2012
	EQ 10MG BASE	A077512 002	Sep 12, 2012
	EQ 20MG BASE	A077512 003	Sep 12, 2012
ESJAY PHARMA	EQ 5MG BASE	A077550 001	May 14, 2015
	EQ 10MG BASE	A077550 002	May 14, 2015
	EQ 20MG BASE	A077550 003	May 14, 2015
HIKMA PHARMS	EQ 5MG BASE	A078766 001	Sep 11, 2012
	EQ 10MG BASE	A078766 002	Sep 11, 2012
	EQ 20MG BASE	A078766 003	Sep 11, 2012
TEVA PHARMS USA	EQ 5MG BASE	A076765 001	Mar 14, 2012
	EQ 10MG BASE	A076765 002	Mar 14, 2012
	EQ 20MG BASE	A076765 003	Mar 14, 2012

ESLICARBAZEPINE ACETATE

TABLET; ORAL

ESLICARBAZEPINE ACETATE

APOTEX	200MG	A211236 001	Dec 07, 2023
	400MG	A211236 002	Dec 07, 2023
	600MG	A211236 003	Dec 07, 2023
	800MG	A211236 004	Dec 07, 2023
LUPIN LTD	200MG	A211246 001	Mar 27, 2024
	400MG	A211246 002	Mar 27, 2024
	600MG	A211246 003	Mar 27, 2024
	800MG	A211246 004	Mar 27, 2024
TORRENT	200MG	A211227 001	Feb 27, 2024
	400MG	A211227 002	Feb 27, 2024
	600MG	A211227 003	Feb 27, 2024
	800MG	A211227 004	Feb 27, 2024

ESMOLOL HYDROCHLORIDE

INJECTABLE; INJECTION

BREVIBLOC

BAXTER HLTHCARE	10MG/ML	N019386 003	Aug 15, 1988
	20MG/ML	N019386 007	May 28, 2003

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ESMOLOL HYDROCHLORIDE

INJECTABLE; INJECTION

ESMOLOL HYDROCHLORIDE

AM REGENT	10MG/ML	A201126	001	Feb 20, 2015
FRESENIUS KABI USA	10MG/ML	A076573	001	May 02, 2005
	20MG/ML	A076573	002	Aug 23, 2023

ESOMEPRAZOLE MAGNESIUM

CAPSULE, DELAYED REL PELLETS; ORAL

ESOMEPRAZOLE MAGNESIUM

AMNEAL PHARMS NY	EQ 20MG BASE	A209647	001	Apr 10, 2019
	EQ 40MG BASE	A209647	002	Apr 10, 2019
GLENMARK SPECLT	EQ 20MG BASE	A209495	001	May 10, 2019
	EQ 40MG BASE	A209495	002	May 10, 2019
HEC PHARM	EQ 20MG BASE	A207265	002	May 18, 2018
	EQ 40MG BASE	A207265	001	May 18, 2018
HETERO LABS LTD III	EQ 20MG BASE	A202784	001	Sep 21, 2015
	EQ 40MG BASE	A202784	002	Sep 21, 2015
SUN PHARM	EQ 20MG BASE	A209735	001	Apr 30, 2018
	EQ 40MG BASE	A209735	002	Apr 30, 2018
TORRENT	EQ 20MG BASE	A203636	001	Oct 19, 2015
	EQ 40MG BASE	A203636	002	Oct 19, 2015

CAPSULE, DELAYED RELEASE; ORAL

ESOMEPRAZOLE MAGNESIUM

HETERO LABS LTD III	EQ 20MG BASE	A208939	001	May 28, 2020
NORVIUM BIOSCIENCE	EQ 20MG BASE	A212376	001	Oct 16, 2019

TABLET, DELAYED RELEASE; ORAL

ESOMEPRAZOLE MAGNESIUM

P AND L	EQ 20MG BASE	A209202	001	Mar 05, 2019
---------	--------------	---------	-----	--------------

TABLET, ORALLY DISINTEGRATING, DELAYED RELEASE; ORAL

ESOMEPRAZOLE MAGNESIUM

+ DEXCEL	EQ 20MG BASE	N214278	001	Oct 20, 2020
----------	--------------	---------	-----	--------------

ESOMEPRAZOLE MAGNESIUM; NAPROXEN

TABLET, DELAYED RELEASE; ORAL

NAPROXEN AND ESOMEPRAZOLE MAGNESIUM

ACTAVIS LABS FL INC	EQ 20MG BASE; 375MG	A204470	001	Aug 24, 2022
	EQ 20MG BASE; 500MG	A204470	002	Aug 24, 2022
MYLAN	EQ 20MG BASE; 375MG	A204920	001	Jul 20, 2021
	EQ 20MG BASE; 500MG	A204920	002	Jul 20, 2021

VIMOVO

+ HORIZON	EQ 20MG BASE; 375MG **	N022511	002	Apr 30, 2010
+	EQ 20MG BASE; 500MG **	N022511	001	Apr 30, 2010

ESOMEPRAZOLE SODIUM

INJECTABLE; INTRAVENOUS

ESOMEPRAZOLE SODIUM

ACCORD HLTHCARE	EQ 40MG BASE/VIAL	A205379	001	Sep 25, 2015
EUGIA PHARMA	EQ 20MG BASE/VIAL	A204657	001	Aug 10, 2016
MYLAN	EQ 20MG BASE/VIAL	A202686	001	May 17, 2017
	EQ 40MG BASE/VIAL	A202686	002	May 17, 2017
SUN PHARM	EQ 20MG BASE/VIAL	A200882	001	Mar 18, 2013
	EQ 40MG BASE/VIAL	A200882	002	Mar 18, 2013

NEXIUM IV

+ ASTRAZENECA	EQ 20MG BASE/VIAL **	N021689	001	Mar 31, 2005
+	EQ 40MG BASE/VIAL	N021689	002	Mar 31, 2005

ESOMEPRAZOLE STRONTIUM

CAPSULE, DELAYED RELEASE; ORAL

ESOMEPRAZOLE STRONTIUM

+ BELCHER	24.65MG	N202342	001	Aug 06, 2013
+	49.3MG	N202342	002	Aug 06, 2013

ESTAZOLAM

TABLET; ORAL

PROSOM

+ ABBOTT	1MG **	N019080	001	Dec 26, 1990
+	2MG **	N019080	002	Dec 26, 1990

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ESTRADIOL

FILM, EXTENDED RELEASE; TRANSDERMAL

ALORA

ABBVIE	0.025MG/24HR	N020655 004	Apr 05, 2002
	0.05MG/24HR	N020655 001	Dec 20, 1996
	0.075MG/24HR	N020655 002	Dec 20, 1996
	0.1MG/24HR	N020655 003	Dec 20, 1996

FEMPATCH

PARKE DAVIS	0.025MG/24HR	N020417 001	Dec 03, 1996
-------------	--------------	-------------	--------------

GEL; TOPICAL

ESTROGEL

ASCEND THERAPS US	0.06%	N021166 001	Feb 09, 2004
-------------------	-------	-------------	--------------

SYSTEM; TRANSDERMAL

ESCLIM

WOMEN FIRST HLTHCARE	0.025MG/24HR	N020847 001	Aug 04, 1998
	0.0375MG/24HR	N020847 002	Aug 04, 1998
	0.05MG/24HR	N020847 003	Aug 04, 1998
	0.075MG/24HR	N020847 004	Aug 04, 1998
	0.1MG/24HR	N020847 005	Aug 04, 1998

ESTRADERM

+ NOVARTIS	0.05MG/24HR	N019081 002	Sep 10, 1986
+	0.1MG/24HR	N019081 003	Sep 10, 1986

ESTRADIOL

ORTHO MCNEIL PHARM	0.05MG/24HR	N021048 001	Sep 20, 1999
	0.075MG/24HR	N021048 002	Sep 20, 1999
	0.1MG/24HR	N021048 003	Sep 20, 1999

VIVELLE

SANDOZ	0.025MG/24HR	N020323 005	Aug 16, 2000
	0.0375MG/24HR	N020323 001	Oct 28, 1994
	0.05MG/24HR	N020323 002	Oct 28, 1994
	0.075MG/24HR	N020323 003	Oct 28, 1994
	0.1MG/24HR	N020323 004	Oct 28, 1994

TABLET; ORAL

ESTRACE

BRISTOL MYERS SQUIBB	0.5MG	A081295 001	Jun 30, 1993
+	1MG	A084499 001	
+	2MG	A084500 001	

ESTRADIOL

DR REDDYS LABS SA	0.5MG	A040114 003	Mar 14, 1996
	1MG	A040114 001	Mar 14, 1996
	2MG	A040114 002	Mar 14, 1996
LANNETT CO INC	0.5MG	A040138 001	Jan 30, 1998
	1MG	A040138 002	Jan 30, 1998
	2MG	A040138 003	Jan 30, 1998
NORVIUM BIOSCIENCE	0.5MG	A040326 001	Apr 21, 1999
	1MG	A040326 002	Apr 21, 1999
	2MG	A040326 003	Apr 21, 1999
USL PHARMA	0.5MG	A040297 001	Apr 17, 2002
	1MG	A040297 002	Apr 17, 2002
	2MG	A040297 003	Apr 17, 2002

GYNODIOL

DURAMED PHARMS BARR	0.5MG	A040212 001	Dec 29, 1997
	1MG	A040212 002	Dec 29, 1997
	1.5MG	A040212 003	Dec 29, 1997
	2MG	A040212 004	Dec 29, 1997

INNOFEM

NOVO NORDISK INC	0.5MG	A040312 001	Nov 19, 1999
	1MG	A040312 002	Nov 19, 1999
	2MG	A040312 003	Nov 19, 1999

TABLET; VAGINAL

VAGIFEM

+ NOVO NORDISK INC	25MCG **	N020908 001	Mar 26, 1999
--------------------	----------	-------------	--------------

ESTRADIOL ACETATE

TABLET; ORAL

FEMTRACE

+ APIL	0.45MG	N021633 001	Aug 20, 2004
+	0.9MG	N021633 002	Aug 20, 2004
+	1.8MG	N021633 003	Aug 20, 2004

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ESTRADIOL CYPIONATE

INJECTABLE; INJECTION

DEPO-ESTRADIOL

PFIZER

1MG/ML

A085470 001

3MG/ML

A085470 002

ESTRADIOL CYPIONATE

DR REDDYS

5MG/ML

A085620 001

ESTRADIOL CYPIONATE; MEDROXYPROGESTERONE ACETATE

INJECTABLE; INTRAMUSCULAR

LUNELLE

+ PHARMACIA AND UPJOHN 5MG/0.5ML; 25MG/0.5ML

N020874 001 Oct 05, 2000

ESTRADIOL CYPIONATE; TESTOSTERONE CYPIONATE

INJECTABLE; INJECTION

DEPO-TESTADIOL

PHARMACIA AND UPJOHN 2MG/ML; 50MG/ML

N017968 001

TESTOSTERONE CYPIONATE-ESTRADIOL CYPIONATE

WATSON LABS

2MG/ML; 50MG/ML

A085603 001 Mar 13, 1986

ESTRADIOL HEMIHYDRATE

EMULSION; TOPICAL

ESTRASORB

+ EXELTIS USA INC 0.25%

N021371 001 Oct 09, 2003

ESTRADIOL VALERATE

INJECTABLE; INJECTION

DELESTROGEN

+ ENDO OPERATIONS 40MG/ML

N009402 003

ESTRADIOL VALERATE

DR REDDYS

20MG/ML

A083547 001

40MG/ML

A083714 001

FOSUN PHARMA

10MG/ML

A040628 001 Oct 04, 2007

20MG/ML

A040628 002 Oct 04, 2007

40MG/ML

A040628 003 Oct 04, 2007

WATSON LABS

10MG/ML

A083546 001

ESTRADIOL VALERATE; TESTOSTERONE ENANTHATE

INJECTABLE; INJECTION

DITATE-DS

SAVAGE LABS 8MG/ML; 180MG/ML

A086423 001

TESTOSTERONE ENANTHATE AND ESTRADIOL VALERATE

WATSON LABS

4MG/ML; 90MG/ML

A085865 001

8MG/ML; 180MG/ML

A085860 001

ESTRADIOL; NORETHINDRONE ACETATE

TABLET; ORAL

ACTIVELLA

+ AMNEAL 0.5MG; 0.1MG **

N020907 002 Dec 28, 2006

ESTRADIOL AND NORETHINDRONE ACETATE

TEVA PHARMS USA

0.5MG; 0.1MG

A200747 001 Mar 08, 2012

ETYQA

AUROBINDO PHARMA LTD 0.5MG; 0.1MG

A214729 001 Jun 30, 2023

1MG; 0.5MG

A214729 002 Jun 30, 2023

ESTRADIOL; NORGESTIMATE

TABLET; ORAL

ESTRADIOL AND NORGESTIMATE

BARR

1MG, 1MG; N/A, 0.09MG

A076812 001 Apr 29, 2005

PREFEST

+ TEVA WOMENS

1MG, 1MG; N/A, 0.09MG **

N021040 001 Oct 22, 1999

ESTRADIOL; PROGESTERONE

CAPSULE; ORAL

ESTRADIOL AND PROGESTERONE

AMNEAL PHARMS

1MG; 100MG

A214293 001 May 16, 2022

ESTRAMUSTINE PHOSPHATE SODIUM

CAPSULE; ORAL

EMCYT

+ PHARMACIA AND UPJOHN EQ 140MG PHOSPHATE

N018045 001

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ESTROGENS, CONJUGATED

TABLET; ORAL

PREMARIN

WYETH PHARMS 2.5MG N004782 002

ESTROGENS, CONJUGATED SYNTHETIC A

CREAM; VAGINAL

SYNTHETIC CONJUGATED ESTROGENS A

TEVA WOMENS 0.625MG/GM N021788 001 Nov 28, 2008

TABLET; ORAL

CENESTIN

+ ASPEN 0.3MG ** N020992 001 Jun 21, 2002

+ 0.45MG ** N020992 005 Feb 05, 2004

+ 0.625MG ** N020992 002 Mar 24, 1999

+ 0.9MG ** N020992 003 Mar 24, 1999

+ 1.25MG ** N020992 004 Mar 13, 2000

ESTROGENS, CONJUGATED SYNTHETIC B

TABLET; ORAL

ENJUVIA

ASPEN 0.3MG N021443 001 Dec 20, 2004

0.45MG N021443 002 Dec 20, 2004

0.625MG ** N021443 003 May 10, 2004

0.9MG N021443 005 Apr 27, 2007

1.25MG ** N021443 004 May 10, 2004

ESTROGENS, CONJUGATED; MEDROXYPROGESTERONE ACETATE

TABLET; ORAL-28

PREMPHASE (PREMARIN; CYCRIN 14/14)

WYETH PHARMS INC 0.625MG, 0.625MG; N/A, 5MG N020303 002 Dec 30, 1994

PREMPRO (PREMARIN; CYCRIN)

WYETH PHARMS INC 0.625MG, 0.625MG; 2.5MG, 2.5MG N020303 001 Dec 30, 1994

ESTROGENS, CONJUGATED; MEPROBAMATE

TABLET; ORAL

MILPREM-200

MEDPOINTE PHARM HLC 0.45MG; 200MG N011045 002

MILPREM-400

MEDPOINTE PHARM HLC 0.45MG; 400MG N011045 001

PMB 200

WYETH AYERST 0.45MG; 200MG N010971 005

PMB 400

WYETH AYERST 0.45MG; 400MG N010971 003

ESTROGENS, ESTERIFIED

TABLET; ORAL

AMNESTROGEN

BRISTOL MYERS SQUIBB 0.3MG A083266 001

0.625MG A083266 002

1.25MG A083266 003

2.5MG A083266 004

ESTERIFIED ESTROGENS

PVT FORM 0.625MG A083414 001

1.25MG A083765 001

2.5MG A085907 001

SANDOZ 1.25MG A085302 001

ESTRATAB

SOLVAY 0.3MG A086715 001

0.625MG A083209 001

1.25MG A083856 001

2.5MG A083857 001

EVEX

ROCHE PALO 0.625MG A084215 001

1.25MG A083376 002

FEMOGEN

PVT FORM 0.625MG A085076 001

1.25MG A085008 001

2.5MG A085007 001

MENEST

MONARCH PHARMS 0.3MG A084951 001

0.625MG A084948 001

1.25MG A084950 001

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ESTROGENS, ESTERIFIED

TABLET; ORAL

MENEST

2.5MG

A084949 001

ESTRONE

INJECTABLE; INJECTION

ESTROGENIC SUBSTANCE

WYETH AYERST

2MG/ML

A083488 001

ESTRONE

DR REDDYS

5MG/ML

A085239 001

WATSON LABS

2MG/ML

A083397 001

NATURAL ESTROGENIC SUBSTANCE-ESTRONE

WATSON LABS

2MG/ML

A085237 001 Nov 23, 1982

THEELIN

PARKEDALE

1MG/ML

N003977 001

2MG/ML

N003977 002

5MG/ML

N003977 003

ESTROPIPATE

CREAM; VAGINAL

OGEN

PHARMACIA AND UPJOHN

1.5MG/GM

A084710 001

TABLET; ORAL

ESTROPIPATE

BARR

0.75MG

A040135 001 Nov 27, 1996

1.5MG

A040135 002 Nov 27, 1996

3MG

A040135 003 Nov 27, 1996

DURAMED PHARMS BARR

0.75MG

A040296 001 Nov 01, 1999

1.5MG

A040296 002 Nov 01, 1999

3MG

A040296 003 Nov 01, 1999

NORVIUM BIOSCIENCE

0.75MG

A040359 001 Aug 26, 1999

1.5MG

A040359 002 Aug 26, 1999

3MG

A040359 003 Aug 26, 1999

WATSON LABS

0.75MG

A081213 001 Sep 23, 1993

1.5MG

A081214 001 Sep 23, 1993

6MG

A081216 001 Sep 23, 1993

WATSON LABS TEVA

3MG

A081215 001 Sep 23, 1993

OGEN .625

+ PFIZER

0.75MG

A083220 001

OGEN 1.25

+ PFIZER

1.5MG

A083220 002

OGEN 2.5

+ PFIZER

3MG

A083220 003

ORTHO-EST

SUN PHARM INDS INC

0.75MG

A089567 001 Feb 27, 1991

1.5MG

A089582 001 Jul 17, 1991

ESZOPICLONE

TABLET; ORAL

ESZOPICLONE

CHARTWELL RX

1MG

A091165 001 Jul 14, 2011

2MG

A091165 002 Jul 14, 2011

3MG

A091165 003 Jul 14, 2011

HIKMA

1MG

A091153 001 Apr 15, 2014

2MG

A091153 002 Apr 15, 2014

3MG

A091153 003 Apr 15, 2014

IPCA LABS LTD

1MG

A206222 001 Dec 29, 2023

2MG

A206222 002 Dec 29, 2023

3MG

A206222 003 Dec 29, 2023

NOSTRUM LABS INC

1MG

A203087 001 May 08, 2019

2MG

A203087 002 May 08, 2019

3MG

A203087 003 May 08, 2019

ETHACRYNIC ACID

TABLET; ORAL

EDECIN

BAUSCH

50MG

N016092 002

ETHACRYNIC ACID

ALVOGEN

25MG

A205709 001 Jul 24, 2018

ENDO OPERATIONS

25MG

A208501 001 Jul 21, 2017

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ETHACRYNIC ACID

TABLET;ORAL

ETHACRYNIC ACID

HIKMA

25MG

A207262 001 Feb 23, 2017

ETHAMBUTOL HYDROCHLORIDE

TABLET;ORAL

ETHAMBUTOL HYDROCHLORIDE

BARR

400MG

A076057 001 Nov 26, 2001

MYAMBUTOL

KANCHAN HLTHCARE

200MG

N016320 002

500MG

N016320 004

ETHCHLORVYNOL

CAPSULE;ORAL

ETHCHLORVYNOL

BANNER PHARMACAPS

100MG

A084463 001

200MG

A084463 002

500MG

A084463 003

750MG

A084463 004

PLACIDYL

ABBVIE

100MG

N010021 004

200MG

N010021 007

500MG

N010021 002

750MG

N010021 010

ETHINAMATE

CAPSULE;ORAL

VALMID

DISTA

500MG

N009750 001

ETHINYL ESTRADIOL

TABLET;ORAL

ESTINYL

SCHERING

0.02MG

N005292 001

0.05MG

N005292 002

0.5MG

N005292 003

FEMINONE

PHARMACIA AND UPJOHN

0.05MG

N016649 001

LYNORAL

ORGANON USA INC

0.01MG

N005490 003

0.05MG

N005490 002

ETHINYL ESTRADIOL; ETHYNODIOL DIACETATE

TABLET;ORAL-21

DEMULEN 1/35-21

GD SEARLE LLC

0.035MG;1MG **

N018168 001

DEMULEN 1/50-21

GD SEARLE LLC

0.05MG;1MG

N016927 001

ZOVIA 1/35E-21

WATSON PHARMS TEVA

0.035MG;1MG

A072720 001 Dec 30, 1991

ZOVIA 1/50E-21

WATSON LABS

0.05MG;1MG

A072722 001 Dec 30, 1991

TABLET;ORAL-28

DEMULEN 1/35-28

GD SEARLE LLC

0.035MG;1MG **

N018160 001

DEMULEN 1/50-28

GD SEARLE LLC

0.05MG;1MG **

N016936 001

ZOVIA 1/35E-28

DR REDDYS LABS SA

0.035MG;1MG

A072721 001 Dec 30, 1991

ETHINYL ESTRADIOL; ETNOGESTREL

RING;VAGINAL

VERARING

DR REDDYS LABS SA

0.015MG/24HR;0.12MG/24HR

A207577 001 Dec 09, 2021

ETHINYL ESTRADIOL; FERROUS FUMARATE; NORETHINDRONE

TABLET;ORAL-28

NORQUEST FE

PFIZER

0.035MG;75MG;1MG

N018926 001 Jul 18, 1986

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ETHINYL ESTRADIOL; FERROUS FUMARATE; NORETHINDRONE ACETATE

TABLET;ORAL-28

NORLESTRIN FE 1/50				
PARKE DAVIS	0.05MG;75MG;1MG		N016766	001
NORLESTRIN FE 2.5/50				
PARKE DAVIS	0.05MG;75MG;2.5MG		N016854	001

ETHINYL ESTRADIOL; LEVONORGESTREL

TABLET;ORAL

FAYOSIM				
LUPIN LTD	0.02MG,0.15MG;0.025MG,0.15MG;0.03MG,0.15MG;0.01MG,N/A		A205943	001 Mar 29, 2016
LEVONORGESTREL AND ETHINYL ESTRADIOL AND ETHINYL ESTRADIOL				
DR REDDYS LABS SA	0.02MG,0.1MG;0.01MG,N/A		A200407	001 Oct 25, 2011
LYBREL				
+ WYETH PHARMS INC	0.02MG;0.09MG **		N021864	001 May 22, 2007
PREVEN EMERGENCY CONTRACEPTIVE KIT				
TEVA BRANDED PHARM	0.05MG;0.25MG		N020946	001 Sep 01, 1998
SYLEVIA				
SUN PHARM	0.03MG;0.15MG		A202988	001 Feb 06, 2019
TABLET;ORAL-21				
ALESSE				
+ CADENCE HEALTH	0.02MG;0.1MG **		N020683	001 Mar 27, 1997
AVIANE-21				
DURAMED PHARMS BARR	0.02MG;0.1MG		A075796	002 Apr 30, 2001
ENPRESSE-21				
DURAMED PHARMS BARR	0.03MG,0.04MG,0.03MG;0.05MG,0.075MG,0.125MG		A075809	001 Jul 16, 2001
LESSINA-21				
BARR	0.02MG;0.1MG		A075803	001 Mar 20, 2002
LEVLITE				
+ BAYER HLTHCARE	0.02MG;0.1MG **		N020860	001 Jul 13, 1998
LEVONORGESTREL AND ETHINYL ESTRADIOL				
BARR	0.02MG;0.1MG		A075862	001 Apr 29, 2003
LEVORA 0.15/30-21				
WATSON LABS	0.03MG;0.15MG		A073592	001 Dec 13, 1993
NORDETTE-21				
TEVA BRANDED PHARM	0.03MG;0.15MG		N018668	001 May 10, 1982
PORTIA-21				
BARR	0.03MG;0.15MG		A075866	001 May 23, 2002
TRIPHASIL-21				
+ WYETH PHARMS	0.03MG,0.04MG,0.03MG;0.05MG,0.075MG,0.125MG **		N019192	001 Nov 01, 1984
TRIVORA-21				
DR REDDYS LABS SA	0.03MG,0.04MG,0.03MG;0.05MG,0.075MG,0.125MG		A074538	001 Dec 18, 1997
TABLET;ORAL-28				
ALESSE				
+ CADENCE HEALTH	0.02MG;0.1MG **		N020683	002 Mar 27, 1997
CERINTA				
SUN PHARM	0.02MG;0.1MG		A202817	001 Jan 07, 2019
ELIFEMME				
XIROMED	0.03MG,0.04MG,0.03MG;0.05MG,0.075MG,0.125MG		A202507	001 Dec 04, 2015
LEVLITE				
+ BAYER HLTHCARE	0.02MG;0.1MG **		N020860	002 Jul 13, 1998
LEVONORGESTREL AND ETHINYL ESTRADIOL				
AMNEAL PHARMS	0.02MG;0.1MG		A201108	001 Feb 05, 2014
	0.03MG;0.15MG		A201095	001 Dec 08, 2014
BARR	0.02MG;0.1MG		A075862	002 Apr 29, 2003
XIROMED	0.02MG;0.1MG		A202247	001 Dec 08, 2014
	0.03MG,0.04MG,0.03MG;0.05MG,0.075MG,0.125MG		A202970	001 Mar 23, 2018
NORDETTE-28				
+ TEVA BRANDED PHARM	0.03MG;0.15MG **		N018782	001 Jul 21, 1982
ORSYTHIA				
ENDO OPERATIONS	0.02MG;0.1MG		A077099	001 May 11, 2011
TRIPHASIL-28				
+ WYETH PHARMS INC	0.03MG,0.04MG,0.03MG;0.05MG,0.075MG,0.125MG **		N019190	001 Nov 01, 1984

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ETHINYL ESTRADIOL; NORELGESTROMIN

FILM, EXTENDED RELEASE; TRANSDERMAL

ORTHO EVRA

+ JANSSEN PHARMS 0.035MG/24HR;0.15MG/24HR ** N021180 001 Nov 20, 2001

ETHINYL ESTRADIOL; NORETHINDRONE

TABLET; ORAL

FEMCON FE

+ APIL 0.035MG;0.4MG ** N021490 001 Nov 14, 2003

TABLET; ORAL-21

BALZIVA-21

BARR 0.035MG;0.4MG A076198 001 Apr 22, 2004

BREVICON 21-DAY

ALLERGAN 0.035MG;0.5MG N017566 001

GENCEPT 10/11-21

BARR 0.035MG,0.035MG;0.5MG,1MG A072694 001 Feb 28, 1992

MODICON 21

ORTHO MCNEIL PHARM 0.035MG;0.5MG ** N017488 001

N.E.E. 1/35 21

LPI 0.035MG;1MG A071541 001 Dec 14, 1987

NORCEPT-E 1/35 21

ORTHO MCNEIL PHARM 0.035MG;1MG A071545 001 Feb 09, 1989

NORETHIN 1/35E-21

WATSON PHARMS TEVA 0.035MG;1MG A071480 001 Apr 12, 1988

NORETHINDRONE AND ETHINYL ESTRADIOL

WATSON LABS 0.035MG;0.4MG A078379 001 Feb 23, 2010

0.035MG;0.5MG A070684 001 Jan 29, 1987

WATSON PHARMS TEVA 0.035MG;1MG A070685 001 Jan 29, 1987

NORETHINDRONE AND ETHINYL ESTRADIOL (10/11)

WATSON LABS 0.035MG,0.035MG;0.5MG,1MG A071043 001 Apr 01, 1988

NORETHINDRONE AND ETHINYL ESTRADIOL (7/14)

WATSON LABS TEVA 0.035MG,0.035MG;0.5MG,1MG A071041 001 Sep 24, 1991

NORINYL 1+35 21-DAY

ALLERGAN 0.035MG;1MG N017565 001

NORTREL 0.5/35-21

BARR 0.035MG;0.5MG A072692 001 Feb 28, 1992

ORTHO-NOVUM 1/35-21

ORTHO MCNEIL PHARM 0.035MG;1MG ** N017489 002

ORTHO-NOVUM 10/11-21

+ ORTHO MCNEIL JANSSEN 0.035MG,0.035MG;0.5MG,1MG ** N018354 001 Jan 11, 1982

ORTHO-NOVUM 7/14-21

ORTHO MCNEIL PHARM 0.035MG,0.035MG;0.5MG,1MG ** N019004 001 Apr 04, 1984

ORTHO-NOVUM 7/7/7-21

JANSSEN PHARMS 0.035MG,0.035MG,0.035MG;0.5MG,0.75MG,1MG N018985 001 Apr 04, 1984

OVCON-35

+ WARNER CHILCOTT 0.035MG;0.4MG ** N018127 001

OVCON-50

WARNER CHILCOTT 0.05MG;1MG N018128 001

TRI-NORINYL 21-DAY

DR REDDYS LABS SA 0.035MG,0.035MG,0.035MG;0.5MG,1MG,0.5MG N018977 001 Apr 13, 1984

TABLET; ORAL-28

BREVICON 28-DAY

ALLERGAN 0.035MG;0.5MG N017743 001

CYCLAFEM 0.5/35

ENDO OPERATIONS 0.035MG;0.5MG A203413 001 Dec 16, 2015

CYCLAFEM 1/35

ENDO OPERATIONS 0.035MG;1MG A076337 001 Nov 12, 2010

CYCLAFEM 7/7/7

ENDO OPERATIONS 0.035MG,0.035MG,0.035MG;0.5MG,0.75MG,1MG A076338 001 Nov 16, 2010

GENCEPT 10/11-28

BARR 0.035MG,0.035MG;0.5MG,1MG A072697 001 Feb 28, 1992

MODICON 28

+ JANSSEN PHARMS 0.035MG;0.5MG N017735 001

N.E.E. 1/35 28

LPI 0.035MG;1MG A071542 001 Dec 14, 1987

NORCEPT-E 1/35 28

ORTHO MCNEIL PHARM 0.035MG;1MG A071546 001 Feb 09, 1989

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ETHINYL ESTRADIOL; NORETHINDRONE

TABLET; ORAL-28

NORETHIN 1/35E-28					
WATSON LABS	0.035MG;1MG	A071481	001	Apr 12, 1988	
NORETHINDRONE AND ETHINYL ESTRADIOL					
DR REDDYS LABS SA	0.035MG;0.5MG	A070686	001	Jan 29, 1987	
MYLAN LABS LTD	0.035MG,0.035MG,0.035MG;0.5MG,0.75MG,1MG	A200486	001	Dec 28, 2015	
	G				
	0.035MG;0.5MG	A200488	001	Oct 21, 2015	
	0.035MG;1MG	A200489	001	Oct 21, 2015	
WATSON LABS	0.035MG,0.035MG,0.035MG;0.5MG,0.75MG,1MG	A076393	001	Feb 04, 2010	
	G				
XIROMED	0.035MG;0.4MG	A200897	001	May 11, 2015	
	0.05MG;1MG	A203006	001	Aug 05, 2013	
NORETHINDRONE AND ETHINYL ESTRADIOL (7/14)					
WATSON LABS	0.035MG,0.035MG;0.5MG,1MG	A071042	001	Sep 24, 1991	
NORINYL 1+35 28-DAY					
ALLERGAN	0.035MG;1MG	N017565	002		
ORTHO-NOVUM 1/35-28					
+ JANSSEN PHARMS	0.035MG;1MG **	N017919	002		
ORTHO-NOVUM 10/11-28					
+ ORTHO MCNEIL JANSSEN	0.035MG,0.035MG;0.5MG,1MG	N018354	002	Jan 11, 1982	
ORTHO-NOVUM 7/14-28					
ORTHO MCNEIL PHARM	0.035MG,0.035MG;0.5MG,1MG **	N019004	002	Apr 04, 1984	
ORTHO-NOVUM 7/7/7-28					
+ JANSSEN PHARMS	0.035MG,0.035MG,0.035MG;0.5MG,0.75MG,1MG	N018985	002	Apr 04, 1984	
	G **				
OVCON-35					
+ WARNER CHILCOTT LLC	0.035MG;0.4MG **	N017716	001		
OVCON-50					
WARNER CHILCOTT LLC	0.05MG;1MG **	N017576	001		
PIRMELLA 1/35					
LUPIN LTD	0.035MG;1MG	A201512	001	Apr 24, 2013	
PIRMELLA 7/7/7					
LUPIN LTD	0.035MG,0.035MG,0.035MG;0.5MG,0.75MG,1MG	A201510	001	Apr 24, 2013	
	G				
RHUZDAH					
AUROBINDO PHARMA	0.035MG;0.4MG	A207585	001	Oct 11, 2022	

ETHINYL ESTRADIOL; NORETHINDRONE ACETATE

CAPSULE; ORAL

NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL AND FERROUS FUMARATE					
AMNEAL PHARMS	0.02MG;1MG	A214292	001	Jul 20, 2021	
DR REDDYS LABS SA	0.02MG;1MG	A213901	001	Apr 28, 2022	

TABLET; ORAL

FEMHRT					
+ APIL	0.0025MG;0.5MG	N021065	001	Jan 14, 2005	
+	0.005MG;1MG **	N021065	002	Oct 15, 1999	
LOESTRIN 24 FE					
+ TEVA BRANDED PHARM	0.02MG;1MG **	N021871	001	Feb 17, 2006	
MINASTRIN 24 FE					
+ APIL	0.02MG;1MG **	N203667	001	May 08, 2013	
NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL					
XIROMED	0.0025MG;0.5MG	A207260	001	Feb 02, 2017	
	0.005MG;1MG	A207259	001	Dec 27, 2016	
NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL AND ETHINYL ESTRADIOL AND FERROUS FUMARATE					
XIROMED	0.01MG,0.01MG;1MG,N/A	A205049	001	May 31, 2016	
NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL AND FERROUS FUMARATE					
AMNEAL PHARMS	0.02MG;1MG	A078267	001	Sep 01, 2009	
	0.02MG;1MG	A207514	001	Sep 11, 2017	
APOTEX	0.02MG;1MG	A208639	001	Mar 21, 2018	
XIROMED	0.02MG;1MG	A202742	001	Oct 30, 2014	
	0.02MG;1MG	A206120	001	Sep 12, 2017	
OSHIH					
AUROBINDO PHARMA	0.02MG;1MG	A216558	001	Dec 06, 2023	
TABLET; ORAL-21					
ESTROSTEP 21					
+ APIL	0.02MG,0.03MG,0.035MG;1MG,1MG,1MG **	N020130	001	Oct 09, 1996	
GILDESS 1.5/30					
ENDO OPERATIONS	0.03MG;1.5MG	A077075	002	Jul 24, 2012	

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ETHINYL ESTRADIOL; NORETHINDRONE ACETATE

TABLET;ORAL-21

GILDESS 1/20				
ENDO OPERATIONS	0.02MG;1MG	A077077	002	Jul 24, 2012
LOESTRIN 21 1.5/30				
+ TEVA BRANDED PHARM	0.03MG;1.5MG	N017875	001	
LOESTRIN 21 1/20				
+ TEVA BRANDED PHARM	0.02MG;1MG **	N017876	001	
LOESTRIN FE 1.5/30				
+ TEVA BRANDED PHARM	0.03MG;1.5MG	N017355	001	
MICROGESTIN 1.5/30				
DR REDDYS LABS SA	0.03MG;1.5MG	A075548	002	Jul 30, 2003
MICROGESTIN 1/20				
DR REDDYS LABS SA	0.02MG;1MG	A075647	002	Jul 30, 2003
NORLESTRIN 21 1/50				
PARKE DAVIS	0.05MG;1MG	N016749	001	
NORLESTRIN 21 2.5/50				
PARKE DAVIS	0.05MG;2.5MG	N016852	001	
TRI-LEGEST 21				
BARR	0.02MG,0.03MG,0.035MG;1MG,1MG,1MG	A076405	001	Oct 26, 2007

TABLET;ORAL-28

ESTROSTEP FE				
+ APIL	0.02MG,0.03MG,0.035MG;1MG,1MG,1MG **	N020130	002	Oct 09, 1996
GILDESS FE 1.5/30				
ENDO OPERATIONS	0.03MG;1.5MG	A077075	001	Apr 28, 2005
GILDESS FE 1/20				
ENDO OPERATIONS	0.02MG;1MG	A077077	001	May 20, 2005
LOESTRIN FE 1/20				
+ TEVA BRANDED PHARM	0.02MG;1MG	N017354	001	
MICROGESTIN FE 1.5/30				
DR REDDYS LABS SA	0.03MG;1.5MG	A075548	001	Feb 05, 2001
MICROGESTIN FE 1/20				
DR REDDYS LABS SA	0.02MG;1MG	A075647	001	Feb 05, 2001
NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL				
DR REDDYS LABS SA	0.02MG,0.03MG,0.035MG;1MG,1MG,1MG	A076629	001	Mar 18, 2010
NORLESTRIN 28 1/50				
PARKE DAVIS	0.05MG;1MG	N016723	001	

TABLET, CHEWABLE, TABLET;ORAL

LO MINASTRIN FE				
+ APIL	0.01MG,0.01MG,N/A;1MG,N/A,N/A	N204654	001	Jul 24, 2013

ETHINYL ESTRADIOL; NORGESTIMATE

TABLET;ORAL-21

ORTHO CYCLEN-21				
JANSSEN PHARMS	0.035MG;0.25MG **	N019653	001	Dec 29, 1989
ORTHO TRI-CYCLEN				
JANSSEN PHARMS	0.035MG,0.035MG,0.035MG;0.18MG,0.215MG,0.25MG **	N019697	002	Jul 03, 1992

TABLET;ORAL-28

NORGESTIMATE AND ETHINYL ESTRADIOL				
AMNEAL PHARMS	0.035MG,0.035MG,0.035MG;0.18MG,0.215MG,0.25MG	A203870	001	Nov 12, 2015
	0.035MG;0.25MG	A203865	001	Oct 27, 2015
MYLAN	0.035MG;0.25MG	A201896	001	Jan 27, 2016
WATSON LABS	0.025MG,0.025MG,0.025MG;0.18MG,0.215MG,0.25MG	A090479	001	Mar 09, 2011
	0.035MG,0.035MG,0.035MG;0.18MG,0.215MG,0.25MG	A076626	001	Aug 17, 2006
	0.035MG;0.25MG	A076627	001	Aug 17, 2006
XIROMED	0.025MG,0.025MG,0.025MG;0.18MG,0.215MG,0.25MG	A202132	001	Sep 09, 2015
	0.035MG,0.035MG,0.035MG;0.18MG,0.215MG,0.25MG	A201897	001	Jan 27, 2016
ORTHO CYCLEN-28				
+ JANSSEN PHARMS	0.035MG;0.25MG **	N019653	002	Dec 29, 1989
ORTHO TRI-CYCLEN				
+ JANSSEN PHARMS	0.035MG,0.035MG,0.035MG;0.18MG,0.215MG,0.25MG **	N019697	001	Jul 03, 1992
ORTHO TRI-CYCLEN LO				
+ JANSSEN PHARMS	0.025MG,0.025MG,0.025MG;0.18MG,0.215MG,0.25MG **	N021241	001	Aug 22, 2002

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ETHINYL ESTRADIOL; NORGESTIMATE

TABLET; ORAL-28

TRI-PREVI-FEM

ENDO OPERATIONS

0.035MG, 0.035MG, 0.035MG; 0.18MG, 0.215MG,
0.25MG

A076335 001 Mar 26, 2004

ETHINYL ESTRADIOL; NORGESTREL

TABLET; ORAL-21

LO/OVRAL

+ CADENCE HEALTH

0.03MG; 0.3MG

N017612 001

LOW-OGESTREL-21

DR REDDYS LABS SA

0.03MG; 0.3MG

A075288 001 Jul 28, 1999

OGESTREL 0.5/50-21

WATSON LABS

0.05MG; 0.5MG **

A075406 001 Dec 15, 1999

OVRAL

WYETH PHARMS

0.05MG; 0.5MG

N016672 001

TABLET; ORAL-28

LO/OVRAL-28

+ WYETH PHARMS

0.03MG; 0.3MG **

N017802 001

NORGESTREL AND ETHINYL ESTRADIOL

MYLAN LABS LTD

0.03MG; 0.3MG

A201828 001 Jun 21, 2016

0.05MG; 0.5MG

A202875 001 May 08, 2017

OGESTREL 0.5/50-28

WATSON LABS

0.05MG; 0.5MG

A075406 002 Dec 15, 1999

OVRAL-28

WYETH PHARMS

0.05MG; 0.5MG

N016806 001

ETHOPROPAZINE HYDROCHLORIDE

TABLET; ORAL

PARSIDOL

PARKE DAVIS

10MG

N009078 003

50MG

N009078 006

100MG

N009078 008

ETHOSUXIMIDE

SYRUP; ORAL

ETHOSUXIMIDE

TEVA PHARMS

250MG/5ML

A081306 001 Jul 30, 1993

ETHOTOIN

TABLET; ORAL

PEGANONE

+ RECORDATI RARE

250MG

N010841 001

500MG

N010841 003

ETHOXZOLAMIDE

TABLET; ORAL

CARDRASE

PHARMACIA AND UPJOHN

62.5MG

N011047 002

125MG

N011047 001

ETHAMIDE

ALLERGAN

125MG

N016144 001

ETHYLESTRENOL

ELIXIR; ORAL

MAXIBOLIN

ORGANON USA INC

2MG/5ML

N014006 002

TABLET; ORAL

MAXIBOLIN

ORGANON USA INC

2MG

N014005 002

ETHYNODIOL DIACETATE; MESTRANOL

TABLET; ORAL-20

OVULEN

GD SEARLE LLC

1MG; 0.1MG

N016029 002

TABLET; ORAL-21

OVULEN-21

GD SEARLE LLC

1MG; 0.1MG

N016029 003

TABLET; ORAL-28

OVULEN-28

GD SEARLE LLC

1MG; 0.1MG

N016705 001

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ETIDOCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

DURANEST

+	ASTRAZENECA	0.5% **	N017751	003	
+		1% **	N017751	005	

ETIDRONATE DISODIUM

INJECTABLE; INJECTION

DIDRONEL

	MGI PHARMA INC	50MG/ML	N019545	001	Apr 20, 1987
--	----------------	---------	---------	-----	--------------

TABLET; ORAL

DIDRONEL

+	APIL	200MG **	N017831	001	
+		400MG **	N017831	002	

ETIDRONATE DISODIUM

	NORVIUM BIOSCIENCE	200MG	A075800	001	Jan 24, 2003
		400MG	A075800	002	Jan 24, 2003

ETODOLAC

CAPSULE; ORAL

ETODOLAC

	ANI PHARMS	200MG	A074840	001	Aug 29, 1997
		200MG	A074844	001	Dec 23, 1997
		200MG	A074899	001	Jul 08, 1997
		300MG	A074840	002	Aug 29, 1997
		300MG	A074844	002	Dec 23, 1997
		300MG	A074899	002	Jul 08, 1997
	BIOPHARM	300MG	A074929	001	Jan 30, 1998
	CHARTWELL MOLECULES	200MG	A074842	001	Jul 17, 1997
		300MG	A074842	002	Jul 17, 1997
	MYLAN	200MG	A075071	001	Sep 30, 1998
		300MG	A075071	002	Sep 30, 1998
	NATCO PHARMA	200MG	A074932	001	May 16, 1997
		300MG	A074932	002	May 16, 1997
	SANDOZ	200MG	A074942	001	Sep 30, 1997
		300MG	A074942	002	Sep 30, 1997

LODINE

+	WYETH PHARMS INC	200MG **	N018922	002	Jan 31, 1991
+		300MG	N018922	003	Jan 31, 1991

TABLET; ORAL

ETODOLAC

	BIOPHARM	400MG	A074927	001	Oct 30, 1997
	CHARTWELL MOLECULES	400MG	A074841	001	Jun 27, 1997
	IVAX SUB TEVA PHARMS	400MG	A074883	001	Feb 28, 1997
		500MG	A074883	002	Nov 20, 1998
	MYLAN	400MG	A075012	001	Sep 30, 1998
		500MG	A075012	002	Sep 30, 1998
	NATCO PHARMA	400MG	A075104	001	Feb 06, 1998
		500MG	A075104	002	Nov 20, 1998
	OXFORD PHARMS	400MG	A074819	001	Feb 28, 1997
		500MG	A074819	002	Apr 28, 1998
	RANBAXY LABS LTD	400MG	A075226	001	Nov 24, 1998
		500MG	A075226	002	Nov 24, 1998
	SHREE HARI INTL	400MG	A074839	001	Jul 11, 1997
		400MG	A074846	001	Feb 28, 1997
	TEVA	400MG	A074847	001	Apr 23, 1999
		400MG	A075009	001	Nov 26, 1997
		500MG	A074847	002	Apr 23, 1999
		500MG	A075009	002	Dec 28, 1999
	WATSON LABS	400MG	A074892	001	Apr 16, 1997
		400MG	A075069	001	Apr 16, 1998
		500MG	A074892	002	Oct 29, 1998

LODINE

+	WYETH PHARMS INC	400MG **	N018922	004	Jul 29, 1993
+		500MG **	N018922	005	Jun 28, 1996

TABLET, EXTENDED RELEASE; ORAL

ETODOLAC

	ACTAVIS ELIZABETH	400MG	A075696	001	Jul 31, 2000
	ANI PHARMS	400MG	A075943	001	Jul 26, 2002
		500MG	A075943	002	Jul 26, 2002

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ETODOLACTABLET, EXTENDED RELEASE;ORAL
ETODOLAC

	600MG	A075943 003	Jul 26, 2002
WATSON LABS FLORIDA	400MG	A075829 001	Nov 30, 2001
	500MG	A075829 002	Nov 30, 2001
LODINE XL			
WYETH PHARMS INC	400MG **	N020584 001	Oct 25, 1996
	500MG **	N020584 003	Jan 20, 1998
+	600MG **	N020584 002	Oct 25, 1996

ETOMIDATE

INJECTABLE; INJECTION

ETOMIDATE

AVET LIFESCIENCES	2MG/ML	A204618 001	Aug 13, 2014
ENDO OPERATIONS	2MG/ML	A091297 001	Jun 20, 2012
LUITPOLD	2MG/ML	A078867 001	Dec 22, 2009
RISING	2MG/ML	A078289 001	Jan 02, 2009

ETONOGESTREL

IMPLANT; IMPLANTATION

IMPLANON

ORGANON	68MG/IMPLANT	N021529 001	Jul 17, 2006
---------	--------------	-------------	--------------

ETOPOSIDE

CAPSULE; ORAL

VEPESID

+	STRIDES SOFTGELS	50MG	N019557 001	Dec 30, 1986
+		100MG	N019557 002	Dec 30, 1986

INJECTABLE; INJECTION

ETOPOSIDE

DASH PHARMS	20MG/ML	A204927 001	Oct 31, 2017
DASH PHARMS NATCO	20MG/ML	A203507 001	Nov 20, 2017
HOSPIRA	20MG/ML	A074320 001	Aug 30, 1995
	20MG/ML	A074351 001	Aug 30, 1995
PHARMACHEMIE BV	20MG/ML	A074227 001	Feb 22, 1996
PIERRE FABRE	20MG/ML	A074813 001	Jul 09, 1997
TEVA PARENTERAL	20MG/ML	A074510 001	Jun 29, 1995
TEVA PHARMS USA	20MG/ML	A074284 001	Feb 10, 1994
WATSON LABS	20MG/ML	A074228 001	Oct 15, 1996
WATSON LABS INC	20MG/ML	A074968 001	Jan 09, 1998

TOPOSAR

TEVA PARENTERAL	20MG/ML	A074166 001	Feb 27, 1995
-----------------	---------	-------------	--------------

VEPESID

+	CORDEN PHARMA	20MG/ML **	N018768 001	Nov 10, 1983
---	---------------	------------	-------------	--------------

ETOPOSIDE PHOSPHATE

INJECTABLE; INJECTION

ETOPOPHOS PRESERVATIVE FREE

BRISTOL MYERS SQUIBB	EQ 500MG BASE/VIAL	N020906 001	Feb 27, 1998
	EQ 1GM BASE/VIAL	N020906 002	Feb 27, 1998

ETRAVIRINE

TABLET; ORAL

ETRAVIRINE

AMNEAL	25MG	A214196 001	Jun 14, 2021
--------	------	-------------	--------------

ETRETINATE

CAPSULE; ORAL

TEGISON

ROCHE	10MG	N019369 001	Sep 30, 1986
	25MG	N019369 002	Sep 30, 1986

EVANS BLUE

INJECTABLE; INJECTION

EVANS BLUE

PARKE DAVIS	0.5% **	N008041 001	
-------------	---------	-------------	--

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

EXEMESTANE

TABLET; ORAL

EXEMESTANE

ALVOGEN	25MG	A200898	001	Jul 28, 2014
AMNEAL PHARMS	25MG	A206421	001	Dec 28, 2018
DR REDDYS LABS SA	25MG	A208764	001	Aug 08, 2019

EXENATIDE SYNTHETIC

FOR SUSPENSION, EXTENDED RELEASE; SUBCUTANEOUS

BYDUREON

+ ASTRAZENECA AB	2MG/VIAL	N022200	001	Jan 27, 2012
------------------	----------	---------	-----	--------------

BYDUREON PEN

+ ASTRAZENECA AB	2MG	N022200	002	Feb 28, 2014
------------------	-----	---------	-----	--------------

EZETIMIBE

TABLET; ORAL

EZETIMIBE

APOTEX	10MG	A208332	001	Jun 12, 2017
RISING	10MG	A201790	001	Apr 26, 2019
TEVA PHARMS USA	10MG	A078724	001	Jun 12, 2017

EZETIMIBE; ROSUVASTATIN CALCIUM

TABLET; ORAL

ROSZET

+ ALTHEA PHARMS	10MG; EQ 5MG BASE	N213072	001	Mar 23, 2021
	10MG; EQ 10MG BASE	N213072	002	Mar 23, 2021
	10MG; EQ 20MG BASE	N213072	003	Mar 23, 2021
	10MG; EQ 40MG BASE	N213072	004	Mar 23, 2021

EZETIMIBE; SIMVASTATIN

TABLET; ORAL

EZETIMIBE AND SIMVASTATIN

ANI PHARMS	10MG; 10MG	A201890	001	Apr 26, 2017
	10MG; 20MG	A201890	002	Apr 26, 2017
	10MG; 40MG	A201890	003	Apr 26, 2017
	10MG; 80MG	A201890	004	Apr 26, 2017

EZOGABINE

TABLET; ORAL

POTIGA

+ GLAXOSMITHKLINE	50MG	N022345	001	Jun 10, 2011
	200MG	N022345	002	Jun 10, 2011
	300MG	N022345	003	Jun 10, 2011
	400MG	N022345	004	Jun 10, 2011

FAMCICLOVIR

TABLET; ORAL

FAMCICLOVIR

CIPLA	125MG	A078278	001	Mar 21, 2011
	250MG	A078278	002	Mar 21, 2011
	500MG	A078278	003	Mar 21, 2011
HIKMA	125MG	A090128	001	Mar 21, 2011
	250MG	A090128	002	Mar 21, 2011
	500MG	A090128	003	Mar 21, 2011
NATCO PHARMA	125MG	A201333	001	Mar 24, 2011
	250MG	A201333	002	Mar 24, 2011
	500MG	A201333	003	Mar 24, 2011
FAMVIR				
+ NOVARTIS	125MG **	N020363	003	Dec 11, 1995
	250MG **	N020363	001	Apr 26, 1996
	500MG **	N020363	002	Jun 29, 1994

FAMOTIDINE

FOR SUSPENSION; ORAL

PEPCID

+ SALIX PHARMS	40MG/5ML **	N019527	001	Feb 02, 1987
----------------	-------------	---------	-----	--------------

INJECTABLE; INJECTION

FAMOTIDINE

APOTEX INC	10MG/ML	A075942	001	Aug 02, 2002
APOTHECON	10MG/ML	A075707	001	Apr 16, 2001
HIKMA	10MG/ML	A075799	001	Apr 30, 2002
HOSPIRA	10MG/ML	A075705	001	Apr 16, 2001
	10MG/ML	A075870	001	Nov 23, 2001

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

FAMOTIDINE

INJECTABLE; INJECTION

FAMOTIDINE					
	10MG/ML		A075905	001	Nov 23, 2001
ZYDUS PHARMS	10MG/ML		A215828	001	Nov 22, 2022
FAMOTIDINE PRESERVATIVE FREE					
APOTHECON	10MG/ML		A075708	001	Apr 16, 2001
EPIC PHARMA LLC	10MG/ML		A076324	001	Nov 27, 2002
HIKMA	10MG/ML		A075789	001	Apr 30, 2002
HOSPIRA	10MG/ML		A075669	001	Apr 16, 2001
FAMOTIDINE PRESERVATIVE FREE (PHARMACY BULK)					
EPIC PHARMA LLC	10MG/ML		A076322	001	Nov 27, 2002
FAMOTIDINE PRESERVATIVE FREE IN PLASTIC CONTAINER					
ABBVIE	0.4MG/ML		A075729	001	Dec 17, 2001
PEPCID					
+ MERCK	10MG/ML **		N019510	001	Nov 04, 1986
PEPCID PRESERVATIVE FREE					
+ MERCK	10MG/ML **		N019510	004	Nov 04, 1986
PEPCID PRESERVATIVE FREE IN PLASTIC CONTAINER					
+ MERCK SHARP DOHME	0.4MG/ML **		N020249	001	Feb 18, 1994

TABLET; ORAL

FAMOTIDINE					
ACTAVIS ELIZABETH	20MG		A075650	001	Sep 14, 2001
	40MG		A075650	002	Sep 14, 2001
APOTEX	10MG		A075610	001	Mar 12, 2002
MYLAN PHARMS INC	20MG		A075457	001	Apr 18, 2001
	40MG		A075457	002	Apr 18, 2001
NATCO PHARMA	10MG		A075674	001	Dec 21, 2001
PERRIGO R AND D	20MG		A077352	002	Jul 27, 2005
	40MG		A077352	001	Jul 27, 2005
RISING	20MG		A075704	001	Apr 16, 2001
	40MG		A075704	002	Apr 16, 2001
SANDOZ	10MG		A076101	001	Oct 21, 2002
	20MG		A075607	001	May 10, 2001
	20MG		A075793	001	Apr 16, 2001
	40MG		A075607	002	May 10, 2001
	40MG		A075793	002	Apr 16, 2001
SUN PHARM INDS LTD	10MG		A090283	001	Nov 17, 2009
	20MG		A090283	002	Nov 17, 2009
SUN PHARM INDUSTRIES	20MG		A075639	002	Dec 12, 2001
	40MG		A075639	001	Dec 12, 2001
TEVA	10MG		A075312	001	May 31, 2001
	20MG		A075311	001	Apr 16, 2001
	40MG		A075311	002	Apr 16, 2001
WATSON LABS	10MG		A075404	001	Nov 28, 2001
	20MG		A075062	002	Apr 16, 2001
	40MG		A075062	001	Apr 16, 2001
PEPCID					
+ BAUSCH	20MG **		N019462	001	Oct 15, 1986
+	40MG **		N019462	002	Oct 15, 1986

TABLET, CHEWABLE; ORAL

FAMOTIDINE					
PERRIGO	10MG		A075715	001	Aug 22, 2003
PEPCID AC					
+ KENVUE BRANDS	10MG **		N020801	001	Sep 24, 1998

TABLET, ORALLY DISINTEGRATING; ORAL

FLUXID					
UCB INC	20MG		N021712	001	Sep 24, 2004
	40MG		N021712	002	Sep 24, 2004
PEPCID RPD					
MERCK	20MG		N020752	001	May 28, 1998
	40MG		N020752	002	May 28, 1998

FAMOTIDINE; IBUPROFEN

TABLET; ORAL

DUEXIS					
+ HORIZON	26.6MG; 800MG **		N022519	001	Apr 23, 2011
FAMOTIDINE AND IBUPROFEN					
TORRENT	26.6MG; 800MG		A215925	001	Mar 18, 2024

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

FEBUXOSTAT

TABLET;ORAL

FEBUXOSTAT

LUPIN LTD	40MG	A205406 001	Jan 17, 2024
	80MG	A205406 002	Jan 17, 2024
MYLAN	40MG	A205385 001	Jul 01, 2019
	80MG	A205385 002	Jul 01, 2019
TORRENT	40MG	A211837 001	Dec 19, 2023
	80MG	A211837 002	Dec 19, 2023

FELODIPINE

TABLET, EXTENDED RELEASE;ORAL

FELODIPINE

JUBILANT GENERICS	2.5MG	A203983 001	Aug 19, 2016
	5MG	A203983 002	Aug 19, 2016
	10MG	A203983 003	Aug 19, 2016
MYLAN	2.5MG	A078855 001	Apr 17, 2008
	5MG	A078855 002	Apr 17, 2008
	10MG	A078855 003	Apr 17, 2008
SUN PHARM INDUSTRIES	2.5MG	A075896 001	Nov 02, 2004
	5MG	A075896 002	Nov 02, 2004
	10MG	A075896 003	Nov 02, 2004
WOCKHARDT	2.5MG	A091484 001	Aug 15, 2012
	5MG	A091484 002	Aug 15, 2012
	10MG	A091484 003	Aug 15, 2012
PLENDIL			
+ ASTRAZENECA	2.5MG **	N019834 004	Sep 22, 1994
+	5MG **	N019834 001	Jul 25, 1991
+	10MG **	N019834 002	Jul 25, 1991

FENOFIBRATE

CAPSULE;ORAL

ANTARA (MICRONIZED)

+ LUPIN	30MG **	N021695 004	Oct 18, 2013
	87MG	N021695 002	Nov 30, 2004
+	90MG **	N021695 005	Oct 18, 2013

FENOFIBRATE (MICRONIZED)

IMPAX LABS	67MG	A075868 001	Oct 27, 2003
	134MG	A075868 002	Oct 27, 2003
	200MG	A075868 003	Oct 27, 2003
INVAGEN PHARMS	67MG	A207378 001	Mar 28, 2017
	134MG	A207378 002	Mar 28, 2017
	200MG	A207378 003	Mar 28, 2017
NORVIUM BIOSCIENCE	43MG	A202579 001	Jan 10, 2013
	130MG	A202579 002	Jan 10, 2013
NOVAST LABS	67MG	A207564 001	Apr 19, 2019
	134MG	A207564 002	Apr 19, 2019
	200MG	A207564 003	Apr 19, 2019
RISING	67MG	A202676 001	Oct 23, 2012
	134MG	A202676 002	Oct 23, 2012
	200MG	A202676 003	Oct 23, 2012

LIPIDIL

ABBVIE	100MG	N019304 001	Dec 31, 1993
--------	-------	-------------	--------------

LIPOFEN

CIPHER PHARMS INC	100MG	N021612 002	Jan 11, 2006
-------------------	-------	-------------	--------------

TRICOR (MICRONIZED)

+ ABBVIE	67MG **	N019304 002	Feb 09, 1998
+	134MG **	N019304 003	Jun 30, 1999
+	200MG **	N019304 004	Jun 30, 1999

TABLET;ORAL

FENOFIBRATE

ALEMBIC	54MG	A213252 001	Jan 17, 2020
	160MG	A213252 002	Jan 17, 2020
MYLAN	107MG	A076520 002	Dec 29, 2005
SUN PHARM INDS LTD	54MG	A076635 001	Oct 31, 2005
	160MG	A076635 003	Oct 31, 2005

FENOGLIDE

+ SALIX	40MG	N022118 001	Aug 10, 2007
+	120MG	N022118 002	Aug 10, 2007

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

FENOFIBRATE

TABLET; ORAL

TRICOR

+	ABBVIE	48MG	N021656	001	Nov 05, 2004
+		54MG **	N021203	001	Sep 04, 2001
+		145MG	N021656	002	Nov 05, 2004
+		160MG **	N021203	003	Sep 04, 2001

TRIGLIDE

SKYEPHARMA AG

		50MG	N021350	001	May 07, 2005
--	--	------	---------	-----	--------------

+		160MG	N021350	002	May 07, 2005
---	--	-------	---------	-----	--------------

FENOFIBRIC ACID

TABLET; ORAL

FIBRICOR

+	ATHENA	35MG	N022418	001	Aug 14, 2009
+		105MG	N022418	002	Aug 14, 2009

FENOLDOPAM MESYLATE

INJECTABLE; INJECTION

CORLOPAM

+	HOSPIRA	EQ 10MG BASE/ML	N019922	001	Sep 23, 1997
---	---------	-----------------	---------	-----	--------------

FENOLDOPAM MESYLATE

HIKMA

LUITPOLD

SANDOZ

TEVA PARENTERAL

		EQ 10MG BASE/ML	A076582	001	Oct 12, 2004
		EQ 10MG BASE/ML	A076656	001	Dec 01, 2003
		EQ 10MG BASE/ML	A077155	001	Feb 15, 2005
		EQ 10MG BASE/ML	A077826	001	Mar 07, 2007

FENOPROFEN CALCIUM

CAPSULE; ORAL

FENOPROFEN CALCIUM

AM THERAP

HALSEY

MISEMER

PAR PHARM

QUANTUM PHARMICS

RISING

WARNER CHILCOTT

WATSON LABS

NALFON

+	KEY THERAP	EQ 200MG BASE	N017604	003	
		EQ 300MG BASE **	N017604	002	
+		EQ 400MG BASE	N017604	004	Jul 21, 2009

TABLET; ORAL

FENOPROFEN CALCIUM

AM THERAP

ANI PHARMS

DAVA PHARMS INC

HALSEY

IVAX SUB TEVA PHARMS

KEY THERAP

QUANTUM PHARMICS

RISING

STRIDES PHARMA

SUN PHARM INDUSTRIES

USL PHARMA

WATSON LABS

WATSON LABS TEVA

		EQ 600MG BASE	A072309	001	Aug 17, 1988
		EQ 600MG BASE	A072274	001	May 02, 1988
		EQ 600MG BASE	A072326	001	Aug 17, 1988
		EQ 600MG BASE	A072357	001	Aug 17, 1988
		EQ 600MG BASE	A072557	001	Aug 29, 1988
		EQ 600MG BASE	A072267	001	Aug 17, 1988
		EQ 600MG BASE	A072194	001	Aug 17, 1988
		EQ 600MG BASE	A072396	001	Oct 17, 1988
		EQ 600MG BASE	A072429	001	Aug 17, 1988
		EQ 600MG BASE	A072902	001	Dec 21, 1990
		EQ 600MG BASE	A072362	001	Aug 17, 1988
		EQ 600MG BASE	A072165	001	Aug 17, 1988
		EQ 600MG BASE	A072602	001	Oct 11, 1988
		EQ 600MG BASE	A072407	001	Aug 17, 1988

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

FENOPROFEN CALCIUM

TABLET; ORAL

NALFON

+ DISTA

EQ 600MG BASE **

N017710 001

FENTANYL

FILM, EXTENDED RELEASE; TRANSDERMAL

DURAGESIC-100

+ JANSSEN PHARMS

100MCG/HR **

N019813 001 Aug 07, 1990

DURAGESIC-12

+ JANSSEN PHARMS

12.5MCG/HR **

N019813 005 Feb 04, 2005

DURAGESIC-25

+ JANSSEN PHARMS

25MCG/HR **

N019813 004 Aug 07, 1990

DURAGESIC-37

+ JANSSEN PHARMS

37.5MCG/HR **

N019813 006 Jan 24, 2018

DURAGESIC-50

+ JANSSEN PHARMS

50MCG/HR **

N019813 003 Aug 07, 1990

DURAGESIC-75

+ JANSSEN PHARMS

75MCG/HR **

N019813 002 Aug 07, 1990

FENTANYL-100

ACTAVIS LABS UT INC

100MCG/HR

A076709 004 Aug 20, 2007

LAVIPHARM LABS

100MCG/HR

A077051 004 Aug 04, 2006

MAYNE PHARMA

100MCG/HR

A077062 004 Aug 20, 2007

NOVEN

100MCG/HR

A077775 004 Oct 16, 2009

ZYDUS PHARMS

100MCG/HR

A209655 008 Jan 24, 2023

FENTANYL-12

ZYDUS PHARMS

12.5MCG/HR

A209655 001 Jan 24, 2023

FENTANYL-25

ACTAVIS LABS UT INC

25MCG/HR

A076709 001 Aug 20, 2007

LAVIPHARM LABS

25MCG/HR

A077051 001 Aug 04, 2006

MAYNE PHARMA

25MCG/HR

A077062 001 Aug 20, 2007

NOVEN

25MCG/HR

A077775 001 Oct 16, 2009

ZYDUS PHARMS

25MCG/HR

A209655 002 Jan 24, 2023

FENTANYL-37

ZYDUS PHARMS

37.5MCG/HR

A209655 003 Jan 24, 2023

FENTANYL-50

ACTAVIS LABS UT INC

50MCG/HR

A076709 002 Aug 20, 2007

LAVIPHARM LABS

50MCG/HR

A077051 002 Aug 04, 2006

MAYNE PHARMA

50MCG/HR

A077062 002 Aug 20, 2007

NOVEN

50MCG/HR

A077775 002 Oct 16, 2009

ZYDUS PHARMS

50MCG/HR

A209655 004 Jan 24, 2023

FENTANYL-62

ZYDUS PHARMS

62.5MCG/HR

A209655 005 Jan 24, 2023

FENTANYL-75

ACTAVIS LABS UT INC

75MCG/HR

A076709 003 Aug 20, 2007

LAVIPHARM LABS

75MCG/HR

A077051 003 Aug 04, 2006

MAYNE PHARMA

75MCG/HR

A077062 003 Aug 20, 2007

NOVEN

75MCG/HR

A077775 003 Oct 16, 2009

ZYDUS PHARMS

75MCG/HR

A209655 006 Jan 24, 2023

FENTANYL-87

ZYDUS PHARMS

87.5MCG/HR

A209655 007 Jan 24, 2023

SPRAY; SUBLINGUAL

SUBSYS

+ BTCP PHARMA

0.1MG **

N202788 001 Jan 04, 2012

+

0.2MG **

N202788 002 Jan 04, 2012

+

0.4MG **

N202788 003 Jan 04, 2012

+

0.6MG **

N202788 004 Jan 04, 2012

+

0.8MG **

N202788 005 Jan 04, 2012

+

1.2MG **

N202788 006 Aug 30, 2012

+

1.6MG **

N202788 007 Aug 30, 2012

FENTANYL CITRATE

FILM; BUCCAL

ONSOLIS

ADALVO

EQ 0.2MG BASE

N022266 001 Jul 16, 2009

EQ 0.4MG BASE

N022266 002 Jul 16, 2009

EQ 0.6MG BASE

N022266 003 Jul 16, 2009

EQ 0.8MG BASE

N022266 004 Jul 16, 2009

EQ 1.2MG BASE

N022266 005 Jul 16, 2009

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

FENTANYL CITRATE

INJECTABLE; INJECTION

FENTANYL CITRATE

ABBOTT	EQ 0.05MG BASE/ML	A070636 001	Apr 30, 1990
	EQ 0.05MG BASE/ML	A070637 001	Apr 30, 1990
WATSON LABS	EQ 0.05MG BASE/ML	A073488 001	Jun 30, 1992
FENTANYL CITRATE PRESERVATIVE FREE			
DR REDDYS	EQ 0.05MG BASE/ML	A074917 001	Feb 03, 1998
HOSPIRA	EQ 0.05MG BASE/ML	A072786 001	Sep 24, 1991

SOLUTION; INTRAVENOUS

FENTANYL CITRATE

+ EXELA PHARMA	EQ 2.5MG BASE/50ML (EQ 0.05MG BASE/ML)	N215870 001	Feb 08, 2023
	**		
+	EQ 5MG BASE/100ML (EQ 0.05MG BASE/ML)	N215870 002	Feb 08, 2023
	**		

SPRAY, METERED; NASAL

LAZANDA

+ BTCP PHARMA	EQ 0.1MG BASE **	N022569 001	Jun 30, 2011
+	EQ 0.3MG BASE **	N022569 003	Dec 21, 2015
+	EQ 0.4MG BASE **	N022569 002	Jun 30, 2011

TABLET; BUCCAL, SUBLINGUAL

FENTANYL CITRATE

DR REDDYS LABS SA	EQ 0.1MG BASE	A206329 001	Aug 22, 2022
	EQ 0.2MG BASE	A206329 002	Aug 22, 2022
	EQ 0.3MG BASE	A206329 003	Aug 22, 2022
	EQ 0.4MG BASE	A206329 004	Aug 22, 2022
	EQ 0.6MG BASE	A206329 005	Aug 22, 2022
	EQ 0.8MG BASE	A206329 006	Aug 22, 2022
WATSON LABS	EQ 0.1MG BASE	A079075 001	Jan 07, 2011
	EQ 0.2MG BASE	A079075 002	Jan 07, 2011
	EQ 0.4MG BASE	A079075 003	Jan 07, 2011
	EQ 0.6MG BASE	A079075 004	Jan 07, 2011
	EQ 0.8MG BASE	A079075 005	Jan 07, 2011

FENTORA

+ CEPHALON	EQ 0.1MG BASE	N021947 001	Sep 25, 2006
+	EQ 0.2MG BASE	N021947 002	Sep 25, 2006
+	EQ 0.3MG BASE **	N021947 006	Mar 02, 2007
+	EQ 0.4MG BASE	N021947 003	Sep 25, 2006
+	EQ 0.6MG BASE	N021947 004	Sep 25, 2006
+	EQ 0.8MG BASE	N021947 005	Sep 25, 2006

TABLET; SUBLINGUAL

ABSTRAL

+ SENTYNL THERAPS INC	EQ 0.1MG BASE **	N022510 001	Jan 07, 2011
+	EQ 0.2MG BASE **	N022510 002	Jan 07, 2011
+	EQ 0.3MG BASE **	N022510 003	Jan 07, 2011
+	EQ 0.4MG BASE **	N022510 004	Jan 07, 2011
+	EQ 0.6MG BASE **	N022510 005	Jan 07, 2011
+	EQ 0.8MG BASE **	N022510 006	Jan 07, 2011

FENTANYL CITRATE

ACTAVIS LABS FL INC	EQ 0.1MG BASE	A207338 001	Nov 17, 2017
	EQ 0.2MG BASE	A207338 002	Nov 17, 2017
	EQ 0.3MG BASE	A207338 003	Nov 17, 2017
	EQ 0.4MG BASE	A207338 004	Nov 17, 2017
	EQ 0.6MG BASE	A207338 005	Nov 17, 2017
	EQ 0.8MG BASE	A207338 006	Nov 17, 2017

TROCHE/LOZENGE; ORAL

FENTANYL

CEPHALON	EQ 0.1MG BASE	N020195 007	Oct 30, 1995
	EQ 0.2MG BASE	N020195 001	Oct 04, 1993
	EQ 0.3MG BASE	N020195 002	Oct 04, 1993
	EQ 0.4MG BASE	N020195 003	Oct 04, 1993

TROCHE/LOZENGE; TRANSMUCOSAL

ACTIQ

+ CEPHALON	EQ 0.2MG BASE	N020747 001	Nov 04, 1998
+	EQ 0.4MG BASE	N020747 002	Nov 04, 1998
+	EQ 0.6MG BASE	N020747 003	Nov 04, 1998
+	EQ 0.8MG BASE	N020747 004	Nov 04, 1998
+	EQ 1.2MG BASE	N020747 005	Nov 04, 1998
+	EQ 1.6MG BASE	N020747 006	Nov 04, 1998

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

FENTANYL CITRATE

TROCHE/LOZENGE; TRANSMUCOSAL

FENTANYL CITRATE

PAR PHARM

EQ 0.2MG BASE

A077312 001 Oct 30, 2009

EQ 0.4MG BASE

A077312 002 Oct 30, 2009

EQ 0.6MG BASE

A077312 003 Oct 30, 2009

EQ 0.8MG BASE

A077312 004 Oct 30, 2009

EQ 1.2MG BASE

A077312 005 Oct 30, 2009

EQ 1.6MG BASE

A077312 006 Oct 30, 2009

SPECGX LLC

EQ 0.2MG BASE

A078907 001 Oct 30, 2009

EQ 0.4MG BASE

A078907 002 Oct 30, 2009

EQ 0.6MG BASE

A078907 003 Oct 30, 2009

EQ 0.8MG BASE

A078907 004 Oct 30, 2009

EQ 1.2MG BASE

A078907 005 Oct 30, 2009

EQ 1.6MG BASE

A078907 006 Oct 30, 2009

FENTANYL HYDROCHLORIDE

SYSTEM; IONTOPHORESIS, TRANSDERMAL

IONSYS

+ THE MEDICINES CO

EQ 40MCG BASE/ACTIVATION

N021338 001 May 22, 2006

FERRIC AMMONIUM CITRATE

FOR SOLUTION; ORAL

FERRISELTZ

OTSUKA

600MG/PACKET

N020292 001 Oct 14, 1997

FERRIC DERISOMALTOSE

SOLUTION; INTRAVENOUS

MONOFERRIC

+ PHARMACOSMOS

100MG/ML (100MG/ML)

N208171 001 Jan 16, 2020

+

500MG/5ML (100MG/ML)

N208171 002 Jan 16, 2020

FERRIC OXYHYDROXIDE

INJECTABLE; INJECTION

DEXFERRUM

AM REGENT

EQ 50MG IRON/ML

N040024 001 Feb 23, 1996

IRON DEXTRAN

SANOFI AVENTIS US

EQ 50MG IRON/ML

N010787 002

PROFERDEX

NEW RIVER

EQ 50MG IRON/ML

N017807 001

INJECTABLE; INTRAVENOUS

VENOFER

AM REGENT

EQ 65MG IRON/3.25ML (EQ 20MG IRON/ML)

N021135 005 Mar 29, 2013

EQ 75MG IRON/3.75ML (EQ 20MG IRON/ML)

N021135 003 Mar 29, 2005

FERRIC PYROPHOSPHATE CITRATE

POWDER; INTRAVENOUS

TRIFERIC

+ ROCKWELL MEDICAL INC

272MG IRON/PACKET

N208551 001 Apr 25, 2016

SOLUTION; INTRAVENOUS

TRIFERIC

+ ROCKWELL MEDICAL INC

27.2MG IRON/5ML (5.44MG IRON/ML)

N206317 001 Jan 23, 2015

+

272MG IRON/50ML (5.44MG IRON/ML)

N206317 002 Sep 04, 2015

TRIFERIC AVNU

+ ROCKWELL MEDICAL INC

6.75MG IRON/4.5ML (1.5MG IRON/ML)

N212860 001 Mar 27, 2020

FERROUS CITRATE, FE-59

INJECTABLE; INJECTION

FERROUS CITRATE FE 59

MALLINCKRODT

25uCi/ML

N016729 001

FERROUS SULFATE; FOLIC ACID

CAPSULE; ORAL

FOLVRON

LEDERLE

182MG; 0.33MG

N006012 003

FERUMOXIDES

INJECTABLE; INJECTION

FERIDEX I.V.

AMAG PHARMS INC

EQ 11.2MG IRON/ML

N020416 001 Aug 30, 1996

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

FERUMOXSIIL

SUSPENSION;ORAL

GASTROMARK

AMAG PHARMS INC

EQ 0.175MG IRON/ML

N020410 001 Dec 06, 1996

FESOTERODINE FUMARATE

TABLET, EXTENDED RELEASE;ORAL

FESOTERODINE FUMARATE

ACCORD HLTHCARE

4MG

A205012 001 Jan 04, 2023

8MG

A205012 002 Jan 04, 2023

ACTAVIS LABS FL INC

4MG

A204868 001 Jan 04, 2023

8MG

A204868 002 Jan 04, 2023

ANI PHARMS

4MG

A204504 001 Jan 04, 2023

8MG

A204504 002 Jan 04, 2023

CHARTWELL RX

4MG

A204983 001 Jan 05, 2023

8MG

A204983 002 Jan 05, 2023

FEXOFENADINE HYDROCHLORIDE

CAPSULE;ORAL

ALLEGRA

CHATTEM SANOFI

60MG **

N020625 001 Jul 25, 1996

FEXOFENADINE HYDROCHLORIDE

BARR

60MG

A076169 001 Jul 13, 2005

SUSPENSION;ORAL

ALLEGRA

+ CHATTEM SANOFI

30MG/5ML

N021963 001 Oct 16, 2006

CHILDREN'S ALLEGRA HIVES

+ CHATTEM SANOFI

30MG/5ML **

N201373 002 Jan 24, 2011

FEXOFENADINE HYDROCHLORIDE

P AND L

30MG/5ML

A201311 001 Jul 25, 2012

TABLET;ORAL

ALLEGRA HIVES

+ CHATTEM SANOFI

60MG

N020872 008 Jan 24, 2011

CHILDREN'S ALLEGRA ALLERGY

+ CHATTEM SANOFI

30MG **

N020872 005 Jan 24, 2011

CHILDREN'S ALLEGRA HIVES

+ CHATTEM SANOFI

30MG

N020872 006 Jan 24, 2011

CHILDREN'S FEXOFENADINE HYDROCHLORIDE ALLERGY

RISING

30MG

A077081 004 Jul 21, 2011

SUN PHARM INDS

30MG

A091567 002 Feb 06, 2012

CHILDREN'S FEXOFENADINE HYDROCHLORIDE HIVES

RISING

30MG

A077081 005 Jul 21, 2011

SUN PHARM INDS

30MG

A091567 001 Feb 06, 2012

FEXOFENADINE HYDROCHLORIDE

BARR

30MG

A076191 001 Aug 31, 2005

60MG

A076191 002 Aug 31, 2005

180MG

A076191 003 Aug 31, 2005

RISING

30MG

A077081 002 Apr 11, 2008

FEXOFENADINE HYDROCHLORIDE ALLERGY

SUN PHARM INDS

60MG

A091567 004 Feb 06, 2012

FEXOFENADINE HYDROCHLORIDE HIVES

SUN PHARM INDS

60MG

A091567 003 Feb 06, 2012

TABLET, ORALLY DISINTEGRATING;ORAL

CHILDREN'S ALLEGRA ALLERGY

+ CHATTEM SANOFI

30MG

N021909 002 Jan 24, 2011

CHILDREN'S ALLEGRA HIVES

+ CHATTEM SANOFI

30MG **

N021909 003 Jan 24, 2011

CHILDREN'S FEXOFENADINE HYDROCHLORIDE ALLERGY

DR REDDYS LABS LTD

30MG

A202978 001 Jan 18, 2013

CHILDREN'S FEXOFENADINE HYDROCHLORIDE HIVES

DR REDDYS LABS LTD

30MG

A202978 002 Jan 18, 2013

FEXOFENADINE HYDROCHLORIDE; PSEUDOEPHEDRINE HYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL

FEXOFENADINE HYDROCHLORIDE AND PSEUDOEPHEDRINE HYDROCHLORIDE

BARR

60MG;120MG

A076236 001 Apr 14, 2005

IMPAX PHARMS

60MG;120MG

A076298 001 Nov 12, 2010

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

FIDAXOMICIN

TABLET; ORAL

FIDAXOMICIN

ACTAVIS LABS FL

200MG

A208443 001 Jan 16, 2024

FINAFLOXACIN

SUSPENSION/DROPS; OTIC

XTORO

+ FONSECA BIOSCIENCES 0.3%

N206307 001 Dec 17, 2014

FINASTERIDE

TABLET; ORAL

FINASTERIDE

ACTAVIS TOTOWA 1MG

A078371 001 Nov 05, 2013

ACTAVIS TOTOWA TEVA 5MG

A077914 001 Mar 28, 2007

CIPLA 1MG

A077335 001 Nov 20, 2014

GEDEON RICHTER USA 5MG

A077251 001 Dec 22, 2006

IVAX SUB TEVA PHARMS 5MG

A076340 001 Jun 19, 2006

MYLAN 5MG

A077578 001 Dec 18, 2006

NATCO PHARMA 1MG

A078161 001 Nov 05, 2013

TEVA 1MG

A076905 001 Nov 05, 2013

FINASTERIDE; TADALAFIL

CAPSULE; ORAL

ENTADFI

+ BLUE WATER BIOTECH 5MG; 5MG

N215423 001 Dec 09, 2021

FINASTERIDE AND TADALAFIL

ZYDUS LIFESCIENCES 5MG; 5MG

A218232 001 Mar 15, 2024

FINGOLIMOD HYDROCHLORIDE

CAPSULE; ORAL

FINGOLIMOD HYDROCHLORIDE

CHARTWELL RX EQ 0.5MG BASE

A207971 001 Jun 29, 2020

SUN PHARM EQ 0.5MG BASE

A208014 001 Dec 04, 2019

TEVA PHARMS USA EQ 0.25MG BASE

A212152 001 Nov 12, 2021

FLAVOXATE HYDROCHLORIDE

TABLET; ORAL

FLAVOXATE HYDROCHLORIDE

IMPAX PHARMS 100MG

A076234 001 Aug 28, 2003

URISPAS

ORTHO MCNEIL JANSSEN 100MG

N016769 001

FLECAINIDE ACETATE

TABLET; ORAL

FLECAINIDE ACETATE

ANI PHARMS 50MG

A076030 001 Oct 28, 2002

100MG

A076030 002 Oct 28, 2002

150MG

A076030 003 Oct 28, 2002

CHARTWELL 50MG

A079164 001 Jul 09, 2009

100MG

A079164 002 Jul 09, 2009

150MG

A079164 003 Jul 09, 2009

TAMBOCOR

+ ALVOGEN 50MG **

N018830 004 Aug 23, 1988

+ 100MG **

N018830 001 Oct 31, 1985

+ 150MG **

N018830 003 Jun 03, 1988

200MG **

N018830 002 Oct 31, 1985

FLORBETAPIR F-18

SOLUTION; INTRAVENOUS

AMYVID

+ AVID RADIOPHARMS INC 10ML (13.5-51mCi/ML)

N202008 001 Apr 06, 2012

+ 10-30ML (13.5-51mCi/ML)

N202008 002 Apr 06, 2012

FLORTAUCIPIR F-18

SOLUTION; INTRAVENOUS

TAUVID

+ AVID RADIOPHARMS INC 30ML (8.1-51mCi/ML)

N212123 001 May 28, 2020

+ 50ML (8.1-51mCi/ML)

N212123 002 May 28, 2020

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

FLOXURIDINE

INJECTABLE; INJECTION

FLOXURIDINE

AM REGENT	50MG/VIAL	A203008	001	Nov 22, 2017
FUDR				
+ HOSPIRA	500MG/VIAL **	N016929	001	

FLUCONAZOLE

FOR SUSPENSION; ORAL

FLUCONAZOLE

HIKMA	50MG/5ML	A076246	001	Jul 29, 2004
	200MG/5ML	A076246	002	Jul 29, 2004
IVAX SUB TEVA PHARMS	50MG/5ML	A077523	001	Sep 12, 2007
	200MG/5ML	A077523	002	Sep 12, 2007
SUN PHARM INDS LTD	50MG/5ML	A076332	001	Jul 29, 2004
	200MG/5ML	A076332	002	Jul 29, 2004
TARO PHARM INDS	50MG/5ML	A076918	001	Dec 18, 2006
	200MG/5ML	A076918	002	Dec 18, 2006

INJECTABLE; INJECTION

DIFLUCAN IN DEXTROSE 5% IN PLASTIC CONTAINER

+ PFIZER	200MG/100ML (2MG/ML)	N019950	003	Sep 29, 1992
+ PFIZER	400MG/200ML (2MG/ML)	N019950	005	Jul 08, 1994

DIFLUCAN IN SODIUM CHLORIDE 0.9%

+ PFIZER	200MG/100ML (2MG/ML)	N019950	001	Jan 29, 1990
+ PFIZER	400MG/200ML (2MG/ML)	N019950	006	Jan 29, 1990

DIFLUCAN IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

+ PFIZER	200MG/100ML (2MG/ML)	N019950	002	Jan 29, 1990
+ PFIZER	400MG/200ML (2MG/ML)	N019950	004	Jan 29, 1990

FLUCONAZOLE IN DEXTROSE 5% IN PLASTIC CONTAINER

HIKMA FARMACEUTICA	200MG/100ML (2MG/ML)	A078764	001	Jan 30, 2012
	400MG/200ML (2MG/ML)	A078764	002	Jan 30, 2012
HOSPIRA	200MG/100ML (2MG/ML)	A076304	001	Jul 29, 2004
	400MG/200ML (2MG/ML)	A076304	002	Jul 29, 2004
NORVIUM BIOSCIENCE	200MG/100ML (2MG/ML)	A076888	001	Mar 25, 2005
	400MG/200ML (2MG/ML)	A076888	002	Mar 25, 2005
WOODWARD	200MG/100ML (2MG/ML)	A077988	001	May 26, 2010
	400MG/200ML (2MG/ML)	A077988	002	May 26, 2010

FLUCONAZOLE IN SODIUM CHLORIDE 0.9%

BAXTER HLTHCARE CORP	200MG/100ML (2MG/ML)	A077947	001	May 26, 2010
	400MG/200ML (2MG/ML)	A077947	002	May 26, 2010
DR REDDYS	200MG/100ML (2MG/ML)	A076653	001	Jul 29, 2004
	400MG/200ML (2MG/ML)	A076653	002	Jul 29, 2004

FLUCONAZOLE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

DR REDDYS	200MG/100ML (2MG/ML)	A076837	001	Jan 13, 2005
	400MG/200ML (2MG/ML)	A076837	002	Jan 13, 2005
HOSPIRA	200MG/100ML (2MG/ML)	A076617	001	Jul 29, 2004
	400MG/200ML (2MG/ML)	A076617	002	Jul 29, 2004
MYLAN LABS LTD	200MG/100ML (2MG/ML)	A076889	001	Mar 25, 2005
	400MG/200ML (2MG/ML)	A076889	002	Mar 25, 2005

TABLET; ORAL

FLUCONAZOLE

ANI PHARMS	50MG	A076086	001	Jul 29, 2004
	100MG	A076086	002	Jul 29, 2004
	150MG	A076086	003	Jul 29, 2004
	200MG	A076086	004	Jul 29, 2004
GEDEON RICHTER USA	50MG	A076432	001	Jul 29, 2004
	100MG	A076432	002	Jul 29, 2004
	150MG	A076432	003	Jul 29, 2004
	200MG	A076432	004	Jul 29, 2004
IVAX SUB TEVA PHARMS	50MG	A076077	001	Jul 29, 2004
	100MG	A076077	002	Jul 29, 2004
	150MG	A076077	003	Jul 29, 2004
	200MG	A076077	004	Jul 29, 2004
LUPIN LTD	50MG	A209146	001	Oct 20, 2023
	100MG	A209146	002	Oct 20, 2023
	150MG	A209146	003	Oct 20, 2023
	200MG	A209146	004	Oct 20, 2023
MYLAN PHARMS INC	50MG	A076042	001	Jul 29, 2004
	100MG	A076042	002	Jul 29, 2004
	150MG	A076042	003	Jul 29, 2004

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

FLUCONAZOLE

TABLET; ORAL

FLUCONAZOLE

	200MG	A076042 004	Jul 29, 2004
NORVIUM BIOSCIENCE	50MG	A076351 001	Jul 29, 2004
	100MG	A076351 002	Jul 29, 2004
	150MG	A076351 003	Jul 29, 2004
	200MG	A076351 004	Jul 29, 2004
PLIVA	50MG	A076424 001	Jul 29, 2004
	100MG	A076424 002	Jul 29, 2004
	150MG	A076424 003	Jul 29, 2004
	200MG	A076424 004	Jul 29, 2004
RANBAXY LABS LTD	50MG	A076386 001	Jul 29, 2004
	100MG	A076386 002	Jul 29, 2004
	150MG	A076386 003	Jul 29, 2004
	200MG	A076386 004	Jul 29, 2004
ROXANE	50MG	A076213 001	Jul 29, 2004
	100MG	A076213 002	Jul 29, 2004
	150MG	A076213 003	Jul 29, 2004
	200MG	A076213 004	Jul 29, 2004
TEVA	50MG	A074681 001	Jul 29, 2004
	100MG	A074681 002	Jul 29, 2004
	150MG	A074681 003	Jul 29, 2004
	200MG	A074681 004	Jul 29, 2004

FLUCYTOSINE

CAPSULE; ORAL

FLUCYTOSINE

HIKMA	250MG	A206550 001	Oct 17, 2017
	500MG	A206550 002	Oct 17, 2017
STRIDES PHARMA	250MG	A207536 001	Jun 18, 2018
	500MG	A207536 002	Jun 18, 2018

FLUDARABINE PHOSPHATE

INJECTABLE; INJECTION

FLUDARA

+ GENZYME CORP	50MG/VIAL **	N020038 001	Apr 18, 1991
FLUDARABINE PHOSPHATE			
ACTAVIS LLC	50MG/2ML (25MG/ML)	A203738 001	Feb 28, 2017
EXTROVIS	50MG/2ML (25MG/ML)	A200647 001	Dec 21, 2011
HOSPIRA	50MG/VIAL	A077790 001	Apr 06, 2007
RISING	50MG/VIAL	A200648 001	Oct 16, 2012
+ SANDOZ	50MG/2ML (25MG/ML) **	N022137 001	Sep 21, 2007

TABLET; ORAL

OFORTA

SANOFI AVENTIS US	10MG	N022273 001	Dec 18, 2008
-------------------	------	-------------	--------------

FLUDEOXYGLUCOSE F-18

INJECTABLE; INJECTION

FLUDEOXYGLUCOSE F18

+ DOWNSTATE CLINCL	4-40mCi/ML **	N020306 001	Aug 19, 1994
+	4-90mCi/ML **	N020306 002	Sep 25, 2001

INJECTABLE; INTRAVENOUS

FLUDEOXYGLUCOSE F18

ESSENTIAL ISOTOPES	20-300mCi/ML	A203946 001	Feb 05, 2014
+ FEINSTEIN	20-200mCi/ML **	N021870 001	Aug 19, 2005
HOT SHOTS NM LLC	4-500mCi/ML	A203937 001	Oct 30, 2014
MIDWEST MEDCL	20-200mCi/ML	A203736 001	Nov 19, 2015
SOFIE	20-500mCi/ML	A203665 001	Feb 14, 2013
UNIV TX SW MEDCTR	20-200mCi/ML	A210265 001	Feb 06, 2020
WEILL MEDCL COLL	10-100mCi/ML **	N021768 001	Aug 05, 2004

FLUDROCORTISONE ACETATE

TABLET; ORAL

FLORINEF

+ CASPER PHARMA LLC	0.1MG **	N010060 001	
FLUDROCORTISONE ACETATE			
CHARTWELL RX	0.1MG	A216013 001	Oct 27, 2022
HIKMA PHARMS	0.1MG	A091302 001	Jul 22, 2011

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

FLUMAZENIL

INJECTABLE; INJECTION

FLUMAZENIL

DR REDDYS	0.5MG/5ML (0.1MG/ML)	A076589 002	Oct 12, 2004
	1MG/10ML (0.1MG/ML)	A076589 001	Oct 12, 2004
RISING	1MG/10ML (0.1MG/ML)	A078595 002	May 13, 2008
ROMAZICON			
+ HOFFMANN LA ROCHE	1MG/10ML (0.1MG/ML) **	N020073 001	Dec 20, 1991
+	0.5MG/5ML (0.1MG/ML) **	N020073 002	Dec 20, 1991

FLUMETHASONE PIVALATE

CREAM; TOPICAL

LOCORTEN

NOVARTIS	0.03%	N016379 001	
----------	-------	-------------	--

FLUNISOLIDE

AEROSOL, METERED; INHALATION

AEROBID

ROCHE PALO	0.25MG/INH	N018340 001	Aug 17, 1984
------------	------------	-------------	--------------

AEROSPAN HEA

+ NORVIUM BIOSCIENCE	0.078MG/INH	N021247 001	Jan 27, 2006
----------------------	-------------	-------------	--------------

SPRAY, METERED; NASAL

FLUNISOLIDE

APOTEX	0.029MG/SPRAY	A077436 001	Aug 09, 2007
--------	---------------	-------------	--------------

NASALIDE

+ IVAX RES	0.025MG/SPRAY **	N018148 001	
------------	------------------	-------------	--

NASAREL

TEVA BRANDED PHARM	0.029MG/SPRAY	N020409 001	Mar 08, 1995
--------------------	---------------	-------------	--------------

FLUOCINOLONE ACETONIDE

CREAM; TOPICAL

FLUCOCET

ALPHARMA US PHARMS	0.025%	A088360 001	Jan 16, 1984
--------------------	--------	-------------	--------------

FLUOCINOLONE ACETONIDE

ALLIED	0.01%	A088047 001	Dec 16, 1982
--------	-------	-------------	--------------

ALPHARMA US PHARMS	0.01%	A088361 001	Jan 16, 1984
--------------------	-------	-------------	--------------

COSETTE	0.025%	A089525 001	Jul 26, 1988
---------	--------	-------------	--------------

INVATECH	0.025%	A088045 001	Dec 16, 1982
----------	--------	-------------	--------------

PERRIGO NEW YORK	0.01%	A086810 001	Mar 04, 1982
------------------	-------	-------------	--------------

	0.025%	A086811 001	Mar 04, 1982
--	--------	-------------	--------------

PHARMAFAIR	0.01%	A088499 001	Aug 02, 1984
------------	-------	-------------	--------------

	0.025%	A088506 001	Aug 02, 1984
--	--------	-------------	--------------

TARO	0.01%	A040035 001	Oct 31, 1994
------	-------	-------------	--------------

	0.01%	A087102 001	Apr 27, 1982
--	-------	-------------	--------------

	0.025%	A040042 001	Oct 31, 1994
--	--------	-------------	--------------

USL PHARMA	0.01%	A088757 001	Feb 11, 1985
------------	-------	-------------	--------------

	0.025%	A088756 001	Mar 28, 1985
--	--------	-------------	--------------

FLUONID

ALLERGAN HERBERT	0.025%	A087156 002	Sep 06, 1984
------------------	--------	-------------	--------------

FLUOTREX

SAVAGE LABS	0.01%	A088174 001	May 06, 1983
-------------	-------	-------------	--------------

	0.025%	A088173 001	Mar 09, 1983
--	--------	-------------	--------------

SYNALAR-HP

MEDIMETRIKS PHARMS	0.2%	N016161 002	
--------------------	------	-------------	--

GEL; TOPICAL

FLUONID

ALLERGAN HERBERT	0.025%	A087300 001	May 27, 1982
------------------	--------	-------------	--------------

OIL; TOPICAL

FLUOCINOLONE ACETONIDE

AMNEAL	0.01%	A201759 001	Oct 17, 2011
--------	-------	-------------	--------------

	0.01%	A201764 001	Oct 17, 2011
--	-------	-------------	--------------

SCIEGEN PHARMS INC	0.01%	A091514 001	Jun 25, 2015
--------------------	-------	-------------	--------------

OIL/DROPS; OTIC

FLUOCINOLONE ACETONIDE

AMNEAL	0.01%	A091306 001	Oct 17, 2011
--------	-------	-------------	--------------

SCIEGEN PHARMS INC	0.01%	A202705 001	Sep 09, 2016
--------------------	-------	-------------	--------------

OINTMENT; TOPICAL

FLUOCINOLONE ACETONIDE

PHARMADERM	0.025%	A088046 001	Dec 16, 1982
------------	--------	-------------	--------------

PHARMAFAIR	0.025%	A088507 001	Feb 27, 1984
------------	--------	-------------	--------------

USL PHARMA	0.025%	A088742 001	Feb 08, 1985
------------	--------	-------------	--------------

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

FLUOCINOLONE ACETONIDE

OINTMENT; TOPICAL

FLUONID

ALLERGAN HERBERT 0.025% A087157 001 Sep 06, 1984

FLUOTREX

SAVAGE LABS 0.025% A088172 001 Mar 09, 1983

SHAMPOO; TOPICAL

CAPEX

+ GALDERMA LABS LP 0.01% N020001 001 Aug 27, 1990

SOLUTION; TOPICAL

FLUOCINOLONE ACETONIDE

ACTAVIS LABS UT INC 0.01% A208386 001 Oct 21, 2016

ALLIED 0.01% A088048 001 Dec 16, 1982

ALPHARMA US PHARMS 0.01% A087159 001 Jun 16, 1982

BAUSCH AND LOMB 0.01% A040059 001 Dec 20, 1993

COSETTE 0.01% A207441 001 Sep 28, 2016

PAI HOLDINGS PHARM 0.01% A088312 001 Jan 27, 1984

PHARMAFAIR 0.01% A088449 001 Feb 08, 1984

FLUONID

ALLERGAN HERBERT 0.01% A087158 001 Mar 17, 1983

FLUOTREX

SAVAGE LABS 0.01% A088171 001 Mar 09, 1983

FLUOCINONIDE

CREAM; TOPICAL

FLUOCINONIDE

AMNEAL 0.1% A211111 001 Jun 04, 2018

ENCUBE 0.05% A211410 001 Oct 16, 2018

PADAGIS ISRAEL 0.1% A090256 001 Jan 14, 2014

PAI HOLDINGS PHARM 0.1% A211758 001 Apr 03, 2019

PERRIGO NEW YORK 0.05% A071790 001 Jul 13, 1988

+ TARO 0.05% ** N019117 001 Jun 26, 1984

LIDEX

+ ALVOGEN 0.05% ** N016908 002

LIDEX-E

+ ALVOGEN 0.05% ** N016908 003

GEL; TOPICAL

FLUOCINONIDE

+ ALVOGEN 0.05% ** N017373 001

COSETTE 0.05% A072537 001 Feb 07, 1989

PADAGIS US 0.05% A209030 001 Jun 19, 2018

OINTMENT; TOPICAL

FLUOCINONIDE

PAI HOLDINGS PHARM 0.05% A207680 001 Sep 28, 2018

LIDEX

+ ALVOGEN 0.05% ** N016909 002

SOLUTION; TOPICAL

FLUOCINONIDE

COSETTE 0.05% A071535 001 Dec 02, 1988

PAI HOLDINGS PHARM 0.05% A207554 001 Mar 18, 2019

TARO 0.05% A072857 001 Aug 02, 1989

TEVA 0.05% A072511 001 Feb 07, 1989

TEVA PHARMS 0.05% A072522 001 Sep 28, 1990

LIDEX

+ ALVOGEN 0.05% ** N018849 001 Apr 06, 1984

FLUORESCEIN SODIUM

INJECTABLE; INJECTION

FUNDUSCEIN-25

+ NOVARTIS 25% ** N017869 001

FLUOROMETHOLONE

CREAM; TOPICAL

OXYLONE

PHARMACIA AND UPJOHN 0.025% N011748 001

OINTMENT; OPHTHALMIC

FML

+ ALLERGAN 0.1% N017760 001 Sep 04, 1985

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

FLUOROMETHOLONE

SUSPENSION/DROPS;OPHTHALMIC

FLUOR-OP

NOVARTIS 0.1% A070185 001 Feb 27, 1986

FLUOROMETHOLONE ACETATE; TOBRAMYCIN

SUSPENSION/DROPS;OPHTHALMIC

TOBRASONE

HARROW EYE 0.1%;0.3% N050628 001 Jul 21, 1989

FLUOROMETHOLONE; SULFACETAMIDE SODIUM

SUSPENSION/DROPS;OPHTHALMIC

FML-S

ALLERGAN 0.1%;10% N019525 001 Sep 29, 1989

FLUOROURACIL

CREAM;TOPICAL

FLUOROPLEX

+ CHARTWELL RX 1% N016988 001

FLUOROURACIL

EXTROVIS 0.5% A203122 001 Apr 20, 2015

INJECTABLE;INJECTION

ADRUCIL

PHARMACIA AND UPJOHN 50MG/ML A081222 001 Jun 28, 1991

+ 50MG/ML ** N017959 001

TEVA PARENTERAL 50MG/ML A040023 001 Oct 18, 1991

50MG/ML A081225 001 Aug 28, 1991

FLUOROURACIL

ABIC 50MG/ML A088929 001 Mar 04, 1986

ABRAXIS PHARM 50MG/ML A089152 001 Mar 21, 1986

50MG/ML A089428 001 Jan 12, 1987

50MG/ML A089519 001 Mar 12, 1987

50MG/ML A089508 001 Jan 26, 1988

BEDFORD 50MG/ML A040772 001 Aug 11, 2008

EBEWE PHARMA 500MG/10ML (50MG/ML) A040291 001 Mar 24, 1999

FRESENIUS KABI USA 50MG/ML A040379 001 Nov 15, 2000

50MG/ML A087791 001 Jan 18, 1983

MARCHAR 50MG/ML A087791 001 Jan 18, 1983

NOVAST LABS 500MG/10ML (50MG/ML) A209219 001 Dec 12, 2019

1GM/20ML (50MG/ML) A209219 002 Dec 12, 2019

2.5GM/50ML (50MG/ML) A209271 002 Dec 11, 2019

5GM/100ML (50MG/ML) A209271 001 Dec 11, 2019

SANDOZ 2.5GM/50ML (50MG/ML) A091299 001 May 02, 2011

5GM/100ML (50MG/ML) A091299 002 May 02, 2011

SMITH AND NEPHEW 50MG/ML A088766 001 Dec 28, 1984

50MG/ML A088767 001 Dec 28, 1984

50MG/ML A089434 001 Mar 26, 1987

SPECTRUM PHARMS 50MG/ML A087792 001 Oct 13, 1982

+ 500MG/10ML (50MG/ML) ** N012209 001

+ 2.5GM/50ML (50MG/ML) ** N012209 002 Jul 29, 2016

TEVA PHARMS USA 500MG/10ML (50MG/ML) A040333 001 Jan 27, 2000

2.5GM/50ML (50MG/ML) A040334 001 Feb 25, 2000

5GM/100ML (50MG/ML) A040334 002 Feb 25, 2000

SOLUTION;TOPICAL

EFUDEX

+ EXTROVIS 5% ** N016831 002

FLUOROPLEX

ELORAC 1% N016765 001

FLUOXETINE HYDROCHLORIDE

CAPSULE;ORAL

FLUOXETINE

SUN PHARM INDUSTRIES EQ 10MG BASE A075787 001 Jan 29, 2002

EQ 20MG BASE A075787 002 Jan 29, 2002

WATSON LABS EQ 10MG BASE A075662 001 Jan 29, 2002

EQ 20MG BASE A075662 002 Jan 29, 2002

FLUOXETINE HYDROCHLORIDE

ACCORD HLTHCARE EQ 10MG BASE A202729 001 Aug 27, 2020

EQ 20MG BASE A202729 002 Aug 27, 2020

EQ 40MG BASE A202729 003 Aug 27, 2020

ANI PHARMS EQ 10MG BASE A076287 001 May 20, 2008

EQ 20MG BASE A076287 002 May 20, 2008

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

FLUOXETINE HYDROCHLORIDE

CAPSULE;ORAL

FLUOXETINE HYDROCHLORIDE

BARR	EQ 10MG BASE	A074803 002	Jan 30, 2002
	EQ 20MG BASE	A074803 001	Aug 02, 2001
	EQ 40MG BASE	A076251 001	May 18, 2005
BEXIMCO PHARMS USA	EQ 10MG BASE	A075807 001	Jan 29, 2002
	EQ 20MG BASE	A075807 002	Jan 29, 2002
CARLSBAD	EQ 10MG BASE	A076022 001	Jan 30, 2002
	EQ 20MG BASE	A076022 002	Jan 30, 2002
CR DOUBLE CRANE	EQ 10MG BASE	A076165 001	Feb 01, 2002
	EQ 20MG BASE	A076165 002	Feb 01, 2002
GRANULES	EQ 10MG BASE	A078143 001	Jan 16, 2008
	EQ 20MG BASE	A078143 002	Jan 16, 2008
	EQ 40MG BASE	A078143 003	Jan 16, 2008
NATCO PHARMA USA	EQ 10MG BASE	A078045 001	Nov 17, 2008
	EQ 20MG BASE	A078045 002	Nov 17, 2008
NORVIUM BIOSCIENCE	EQ 10MG BASE	A075207 001	Jan 30, 2002
	EQ 10MG BASE	A075577 001	Jan 29, 2002
	EQ 20MG BASE	A075207 002	Jan 30, 2002
	EQ 20MG BASE	A075577 002	Jan 29, 2002
	EQ 40MG BASE	A075207 003	May 25, 2007
SANDOZ	EQ 10MG BASE	A077469 001	Nov 17, 2008
	EQ 20MG BASE	A077469 002	Nov 17, 2008
SPECGX LLC	EQ 10MG BASE	A075658 001	Jan 29, 2002
	EQ 20MG BASE	A075658 002	Jan 29, 2002
SUN PHARM INDS LTD	EQ 40MG BASE	A076990 001	Dec 13, 2004

PROZAC

ELI LILLY AND CO	EQ 60MG BASE	N018936 004	Jun 15, 1999
------------------	--------------	-------------	--------------

SARAFEM

+ ELI LILLY AND CO	EQ 10MG BASE **	N018936 007	Jul 06, 2000
+	EQ 20MG BASE **	N018936 008	Jul 06, 2000

CAPSULE, DELAYED REL PELLETS;ORAL

FLUOXETINE HYDROCHLORIDE

BARR	EQ 90MG BASE	A076237 001	Mar 24, 2010
PROZAC WEEKLY			
+ LILLY	EQ 90MG BASE **	N021235 001	Feb 26, 2001

SOLUTION;ORAL

FLUOXETINE HYDROCHLORIDE

ACTAVIS MID ATLANTIC	EQ 20MG BASE/5ML	A075690 001	Jan 31, 2002
BAJAJ	EQ 20MG BASE/5ML	A075920 001	Jan 29, 2002
CHARTWELL MOLECULAR	EQ 20MG BASE/5ML	A076458 001	May 14, 2004
PHARMOBEDIANT CNSLTG	EQ 20MG BASE/5ML	A075514 001	Aug 29, 2002
SCIEGEN PHARMS INC	EQ 20MG BASE/5ML	A075525 001	Jun 27, 2002

PROZAC

+ LILLY	EQ 20MG BASE/5ML **	N020101 001	Apr 24, 1991
---------	---------------------	-------------	--------------

TABLET;ORAL

FLUOXETINE HYDROCHLORIDE

ADAPTIS	EQ 10MG BASE	A211444 001	Sep 13, 2022
	EQ 20MG BASE	A211444 002	Sep 13, 2022
BARR	EQ 10MG BASE	A075810 001	Feb 01, 2002
FOSUN PHARMA	EQ 10MG BASE	A076024 001	Jan 29, 2002
G AND W LABS INC	EQ 60MG BASE	A212191 001	Jul 05, 2019
IVAX SUB TEVA PHARMS	EQ 10MG BASE	A075865 001	Feb 28, 2002
	EQ 40MG BASE	A075865 003	Aug 30, 2004
RISING	EQ 10MG BASE	A075755 001	Aug 02, 2001
	EQ 20MG BASE	A075755 002	Aug 02, 2001
TEVA	EQ 10MG BASE	A075872 001	Jan 29, 2002
	EQ 20MG BASE	A075872 002	Jan 04, 2019
TEVA PHARMS USA	EQ 60MG BASE	A211051 001	Dec 03, 2018
UPSHER SMITH LABS	EQ 10MG BASE	A211696 001	Jan 30, 2019
	EQ 20MG BASE	A211696 002	Jan 30, 2019

PROZAC

+ LILLY	EQ 10MG BASE **	N020974 001	Mar 09, 1999
+	EQ 20MG BASE **	N020974 002	Mar 09, 1999

SARAFEM

+ APIL	EQ 10MG BASE **	N021860 001	May 19, 2006
+	EQ 15MG BASE **	N021860 002	May 19, 2006
+	EQ 20MG BASE **	N021860 003	May 19, 2006

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

FLUOXETINE HYDROCHLORIDE

TABLET; ORAL

SELFEMRA

TEVA PHARMS USA	EQ 10MG BASE	A200151 001	Feb 03, 2014
	EQ 15MG BASE	A200151 002	Feb 03, 2014
	EQ 20MG BASE	A200151 003	Feb 03, 2014

FLUOXYMESTERONE

TABLET; ORAL

ANDROID-F

VALEANT PHARM INTL	10MG	A087196 001	
FLUOXYMESTERONE			
UPSHER SMITH LABS	10MG	A088342 001	Oct 21, 1983
VALEANT PHARM INTL	10MG	A088221 001	May 05, 1983
WATSON LABS	2MG	A088260 001	Dec 06, 1983
	5MG	A088265 001	Dec 06, 1983
	10MG	A088309 001	Dec 06, 1983

HALOTESTIN

PHARMACIA AND UPJOHN	2MG	N010611 002	
	5MG	N010611 006	
	10MG	N010611 010	

ORA-TESTRYL

BRISTOL MYERS SQUIBB	2MG	N011359 001	
	5MG	N011359 002	

FLUPHENAZINE DECANOATE

INJECTABLE; INJECTION

FLUPHENAZINE DECANOATE

HOSPIRA	25MG/ML	A074966 001	Apr 16, 1998
TEVA PARENTERAL	25MG/ML	A074795 001	Sep 10, 1996
PROLIXIN DECANOATE			
+ BRISTOL MYERS SQUIBB	25MG/ML **	N016727 001	

FLUPHENAZINE ENANTHATE

INJECTABLE; INJECTION

PROLIXIN ENANTHATE

APOTHECON	25MG/ML **	N016110 001	
-----------	------------	-------------	--

FLUPHENAZINE HYDROCHLORIDE

CONCENTRATE; ORAL

FLUPHENAZINE HYDROCHLORIDE

ANI PHARMS	5MG/ML	A073058 001	Aug 30, 1991
PERMITIL			
SCHERING	5MG/ML **	N016008 001	
PROLIXIN			
APOTHECON	5MG/ML	A070533 001	Nov 07, 1985

ELIXIR; ORAL

FLUPHENAZINE HYDROCHLORIDE

ANI PHARMS	2.5MG/5ML	A081310 001	Apr 29, 1993
PROLIXIN			
+ APOTHECON	2.5MG/5ML **	N012145 003	

INJECTABLE; INJECTION

PROLIXIN

APOTHECON	2.5MG/ML **	N011751 005	
-----------	-------------	-------------	--

TABLET; ORAL

FLUPHENAZINE HYDROCHLORIDE

PRASCO	1MG	A089804 002	Aug 12, 1988
	2.5MG	A089804 003	Aug 12, 1988
	5MG	A089804 004	Aug 12, 1988
	10MG	A089804 001	Aug 12, 1988
SANDOZ	1MG	A089586 002	Oct 16, 1987
	2.5MG	A089586 003	Oct 16, 1987
	5MG	A089586 004	Oct 16, 1987
	10MG	A089586 001	Oct 16, 1987
TORRENT	1MG	A217094 001	Jan 22, 2024
	2.5MG	A217094 002	Jan 22, 2024
	5MG	A217094 003	Jan 22, 2024
	10MG	A217094 004	Jan 22, 2024
WATSON LABS	1MG	A088555 001	Dec 18, 1987
	2.5MG	A088544 001	Dec 18, 1987
	5MG	A088527 001	Dec 18, 1987

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

FLUPHENAZINE HYDROCHLORIDE

TABLET;ORAL

FLUPHENAZINE HYDROCHLORIDE

	10MG	A088550 001	Dec 18, 1987
PERMITIL			
SCHERING	0.25MG	N012034 001	
	2.5MG	N012034 004	
	5MG	N012034 005	
	10MG	N012034 006	
PROLIXIN			
+ APOTHECON	1MG **	N011751 004	
+	2.5MG **	N011751 001	
+	5MG **	N011751 003	
+	10MG **	N011751 002	

TABLET, EXTENDED RELEASE;ORAL

PERMITIL

SCHERING	1MG	N012419 004	
----------	-----	-------------	--

FLUPREDNISOLONE

TABLET;ORAL

ALPHADROL

PHARMACIA AND UPJOHN	1.5MG	N012259 002	
----------------------	-------	-------------	--

FLURANDRENOLIDE

CREAM;TOPICAL

CORDRAN SP

+ ALMIRALL	0.025% **	N012806 003	
+	0.05%	N012806 002	

FLURANDRENOLIDE

CHARTWELL RX	0.05%	A205342 001	Apr 13, 2016
--------------	-------	-------------	--------------

LOTION;TOPICAL

CORDRAN

+ ALMIRALL	0.05%	N013790 001	
------------	-------	-------------	--

FLURANDRENOLIDE

ALPHARMA US PHARMS	0.05%	A087203 001	Apr 29, 1982
CHARTWELL RX	0.05%	A205343 001	Dec 22, 2016

OINTMENT;TOPICAL

CORDRAN

+ ALMIRALL	0.025% **	N012806 004	
+	0.05%	N012806 001	

FLURANDRENOLIDE; NEOMYCIN SULFATE

CREAM;TOPICAL

CORDRAN N

LILLY	0.05%;EQ 3.5MG BASE/GM	N050346 001	
-------	------------------------	-------------	--

OINTMENT;TOPICAL

CORDRAN N

LILLY	0.05%;EQ 3.5MG BASE/GM	N050345 001	
-------	------------------------	-------------	--

FLURAZEPAM HYDROCHLORIDE

CAPSULE;ORAL

DALMANE

VALEANT PHARM INTL	15MG **	N016721 001	
+	30MG **	N016721 002	

FLURAZEPAM HYDROCHLORIDE

CHARTWELL RX	15MG	A071205 001	Nov 25, 1986
	30MG	A071068 001	Nov 25, 1986
ESJAY PHARMA	15MG	A070444 001	Mar 20, 1986
	30MG	A070445 001	Mar 20, 1986
HALSEY	15MG	A071808 001	Jan 07, 1988
	30MG	A071809 001	Jan 07, 1988
HIKMA	30MG	A071108 001	Dec 08, 1986
HIKMA INTL PHARMS	15MG	A071107 001	Dec 08, 1986
PUREPAC PHARM	15MG	A071927 001	Sep 09, 1987
	30MG	A071551 001	Sep 09, 1987
RISING	15MG	A071717 002	Jul 31, 1991
	30MG	A071717 001	Jul 31, 1991
SUN PHARM INDUSTRIES	15MG	A070454 001	Aug 04, 1986
	30MG	A070455 001	Aug 04, 1986
SUPERPHARM	15MG	A071659 001	Aug 04, 1988
	30MG	A071660 001	Aug 04, 1988

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

FLURAZEPAM HYDROCHLORIDE

CAPSULE;ORAL

FLURAZEPAM HYDROCHLORIDE

USL PHARMA	15MG	A070562	001	Jul 09, 1987
	30MG	A070563	001	Jul 09, 1987
WARNER CHILCOTT	15MG	A071767	001	Dec 04, 1987
	30MG	A071768	001	Dec 04, 1987

FLURBIPROFEN

TABLET;ORAL

ANSAID

+ PHARMACIA AND UPJOHN	50MG	N018766	002	Oct 31, 1988
+	100MG	N018766	003	Oct 31, 1988

FLURBIPROFEN

IVAX SUB TEVA PHARMS	50MG	A074411	001	May 31, 1995
	100MG	A074411	002	May 31, 1995
NATCO PHARMA	50MG	A074358	001	Jun 20, 1994
	100MG	A074358	002	Jun 20, 1994
PLIVA	50MG	A074647	001	Apr 01, 1997
	100MG	A074647	002	Apr 01, 1997
RISING	50MG	A074448	001	Jul 28, 1995
	100MG	A074448	002	Jul 28, 1995
SUN PHARM INDS INC	50MG	A075058	001	Apr 27, 2001
	100MG	A075058	002	Apr 27, 2001
TEVA	50MG	A074405	002	May 24, 1995
	100MG	A074405	001	May 24, 1995
THERAGEN	100MG	A074560	002	May 16, 1997

FLURBIPROFEN SODIUM

SOLUTION/DROPS;OPHTHALMIC

OCUFEN

+ ALLERGAN	0.03% **	N019404	001	Dec 31, 1986
------------	----------	---------	-----	--------------

FLUTAMIDE

CAPSULE;ORAL

EULEXIN

+ SCHERING	125MG **	N018554	001	Jan 27, 1989
------------	----------	---------	-----	--------------

FLUTAMIDE

ACTAVIS LABS FL INC	125MG	A075820	001	Sep 18, 2001
CHARTWELL RX	125MG	A075818	001	Sep 18, 2001
CIPLA	125MG	A075780	001	Sep 19, 2001
NORVIUM BIOSCIENCE	125MG	A076224	001	May 09, 2003

FLUTEMETAMOL F-18

INJECTABLE;INTRAVENOUS

VIZAMYL

+ GE HEALTHCARE	40.5mCi/10ML (4.05mCi/ML)	N203137	001	Oct 25, 2013
-----------------	---------------------------	---------	-----	--------------

FLUTICASONE PROPIONATE

AEROSOL, METERED;INHALATION

FLOVENT

GLAXOSMITHKLINE	0.044MG/INH	N020548	001	Mar 27, 1996
	0.11MG/INH	N020548	002	Mar 27, 1996
	0.22MG/INH	N020548	003	Mar 27, 1996

CREAM;TOPICAL

CUTIVATE

+ FOUGERA PHARMS	0.05% **	N019958	001	Dec 18, 1990
------------------	----------	---------	-----	--------------

FLUTICASONE PROPIONATE

ENCUBE	0.05%	A076633	001	May 14, 2004
NESHER PHARMS	0.05%	A076865	001	Sep 10, 2004

LOTION;TOPICAL

CUTIVATE

+ FOUGERA PHARMS	0.05% **	N021152	001	Mar 31, 2005
------------------	----------	---------	-----	--------------

OINTMENT;TOPICAL

CUTIVATE

+ FOUGERA PHARMS	0.005% **	N019957	001	Dec 14, 1990
------------------	-----------	---------	-----	--------------

FLUTICASONE PROPIONATE

BRIGHT	0.005%	A215343	001	Sep 01, 2023
FOUGERA PHARMS	0.005%	A076300	001	May 14, 2004
TARO PHARM INDS	0.005%	A077145	001	Jun 14, 2005

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

FLUTICASONE PROPIONATE

POWDER; INHALATION

ARMONAIR DIGIHALER

+	TEVA PHARM	0.03MG/INH	N208798	008	Apr 08, 2022
+		0.055MG/INH	N208798	004	Feb 20, 2020
+		0.113MG/INH	N208798	005	Feb 20, 2020
+		0.232MG/INH	N208798	006	Feb 20, 2020

ARMONAIR RESPICLICK

+	TEVA PHARM	0.03MG/INH	N208798	007	Jul 09, 2021
+		0.055MG/INH	N208798	001	Jan 27, 2017
+		0.113MG/INH	N208798	002	Jan 27, 2017
+		0.232MG/INH	N208798	003	Jan 27, 2017

FLOVENT

	GLAXOSMITHKLINE	0.044MG/INH	N020549	001	Nov 07, 1997
		0.088MG/INH	N020549	002	Nov 07, 1997
		0.22MG/INH	N020549	003	Nov 07, 1997

SPRAY, METERED; NASAL

FLONASE

+	HALEON US HOLDINGS	0.05MG/SPRAY **	N020121	001	Oct 19, 1994
---	--------------------	-----------------	---------	-----	--------------

FLUTICASONE PROPIONATE

	HIKMA	0.05MG/SPRAY	A077570	001	Jan 16, 2008
		0.05MG/SPRAY	A208024	001	Apr 17, 2019

FLUTICASONE PROPIONATE; SALMETEROL XINAFOATE

POWDER; INHALATION

AIRDUO DIGIHALER

+	TEVA PHARM	0.055MG/INH;EQ 0.014MG BASE/INH	N208799	004	Jul 12, 2019
+		0.113MG/INH;EQ 0.014MG BASE/INH	N208799	005	Jul 12, 2019
+		0.232MG/INH;EQ 0.014MG BASE/INH	N208799	006	Jul 12, 2019

FLUVASTATIN SODIUM

CAPSULE; ORAL

LESCOL

+	NOVARTIS	EQ 20MG BASE **	N020261	001	Dec 31, 1993
+		EQ 40MG BASE **	N020261	002	Dec 31, 1993

TABLET, EXTENDED RELEASE; ORAL

FLUVASTATIN SODIUM

	NORVIUM BIOSCIENCE	EQ 80MG BASE	A202458	001	Sep 11, 2015
--	--------------------	--------------	---------	-----	--------------

FLUVOXAMINE MALEATE

CAPSULE, EXTENDED RELEASE; ORAL

FLUVOXAMINE MALEATE

	TORRENT	100MG	A203240	001	Oct 31, 2014
		150MG	A203240	002	Oct 31, 2014

LUVOX CR

+	JAZZ PHARMS	100MG **	N022033	001	Feb 28, 2008
+		150MG **	N022033	002	Feb 28, 2008

TABLET; ORAL

FLUVOXAMINE MALEATE

	ACTAVIS ELIZABETH	25MG	A075901	001	Dec 28, 2000
		50MG	A075901	002	Dec 28, 2000
		100MG	A075901	003	Dec 28, 2000
	ANI PHARMS	25MG	A075897	001	Jan 25, 2001
		25MG	A075898	001	Mar 12, 2001
		50MG	A075897	002	Jan 25, 2001
		50MG	A075898	002	Mar 12, 2001
		100MG	A075897	003	Jan 25, 2001
		100MG	A075898	003	Mar 12, 2001
	CHARTWELL RX	25MG	A075900	001	Feb 23, 2006
		50MG	A075900	002	Feb 23, 2006
		100MG	A075900	003	Feb 23, 2006
	HERITAGE PHARMA	25MG	A075894	001	Apr 18, 2001
		50MG	A075894	002	Apr 18, 2001
		100MG	A075894	003	Apr 18, 2001
	MYLAN	50MG	A075950	001	Oct 15, 2001
		100MG	A075950	002	Oct 15, 2001
	NORVIUM BIOSCIENCE	25MG	A075889	001	Nov 29, 2000
		50MG	A075889	002	Nov 29, 2000
		100MG	A075889	003	Nov 29, 2000
	SUN PHARM INDUSTRIES	25MG	A076125	001	Apr 29, 2002
		50MG	A076125	002	Apr 29, 2002

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

FLUVOXAMINE MALEATE

TABLET; ORAL

FLUVOXAMINE MALEATE

	100MG	A076125 003	Apr 29, 2002
SYNTHON PHARMS	25MG	A075899 001	Jan 17, 2001
	50MG	A075899 002	Jan 17, 2001
	100MG	A075899 003	Jan 17, 2001
TEVA	25MG	A075893 001	Sep 10, 2002
	50MG	A075893 002	Sep 10, 2002
	100MG	A075893 003	Sep 10, 2002
UPSHER SMITH LABS	25MG	A075887 001	Jan 05, 2001
	50MG	A075887 002	Jan 05, 2001
	100MG	A075887 003	Jan 05, 2001
LUVOX			
+ SOLVAY	25MG **	N020243 001	Dec 05, 1994
+	50MG **	N020243 002	Dec 05, 1994
+	100MG **	N020243 003	Dec 05, 1994
+	150MG **	N020243 004	Dec 05, 1994

FOLIC ACID

INJECTABLE; INJECTION

FOLIC ACID

BEN VENUE	5MG/ML	A081066 001	Dec 29, 1993
-----------	--------	-------------	--------------

FOLVITE

+ WYETH PHARMS INC	5MG/ML **	N005897 008	
--------------------	-----------	-------------	--

TABLET; ORAL

FOLIC ACID

BARR	1MG	A089177 001	Jan 08, 1986
CONTRACT PHARMACAL	1MG	A085061 001	
ENDO OPERATIONS	1MG	A040756 001	Jun 04, 2010
EVERYLIFE	1MG	A080755 001	
HALSEY	1MG	A083598 001	
HIKMA PHARMS	1MG	A080600 001	
IMPAX LABS	1MG	A080686 001	
IVAX SUB TEVA PHARMS	1MG	A083000 001	
JUBILANT CADISTA	1MG	A040514 001	Jun 14, 2005
LANNETT	1MG	A080816 001	
LILLY	1MG **	N006135 003	
MK LABS	1MG	A083526 001	
NEXGEN PHARMA INC	1MG	A084915 001	
PHARMERAL	1MG	A084158 001	
PIONEER PHARMS	1MG	A088949 001	Sep 13, 1985
PUREPAC PHARM	1MG	A080784 001	
SANDOZ	1MG	A084472 001	
SUN PHARM INDUSTRIES	1MG	A040582 001	Jul 18, 2005
TABLICAPS	1MG	A083133 002	
UDL	1MG	A088199 001	Mar 29, 1983
USL PHARMA	1MG	A087828 001	May 13, 1982
VALEANT PHARM INTL	1MG	A080903 001	
VANGARD	1MG	A088730 001	Mar 23, 1984
VINTAGE PHARMS	1MG	A086296 001	
WATSON LABS	1MG	A083141 001	
	1MG	A085141 002	
WHITEWORTH TOWN PLSN	1MG	A080691 002	
FOLICET			
MISSION PHARMA	1MG	A087438 001	
FOLVITE			
WYETH PHARMS INC	1MG **	N005897 004	

FOMEPIZOLE

INJECTABLE; INJECTION

ANTIZOL

+ PAR PHARM INC	1.5GM/1.5ML (1GM/ML) **	N020696 001	Dec 04, 1997
-----------------	-------------------------	-------------	--------------

FOMEPIZOLE

MYLAN INSTITUTIONAL	1.5GM/1.5ML (1GM/ML)	A079033 001	Apr 07, 2009
---------------------	----------------------	-------------	--------------

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

FOMIVIRSEN SODIUM

INJECTABLE; INJECTION

VITRAVENE PRESERVATIVE FREE

NOVARTIS

6.6MG/ML

N020961 001 Aug 26, 1998

FORMOTEROL FUMARATE

POWDER; INHALATION

FORADIL

+ NOVARTIS

0.012MG/INH **

N020831 001 Feb 16, 2001

FORADIL CERTIHALER

NOVARTIS

0.0085MG/INH

N021592 001 Dec 15, 2006

FOSAMPRENAVIR CALCIUM

SUSPENSION; ORAL

LEXIVA

+ VIIV HLTHCARE

EQ 50MG BASE/ML

N022116 001 Jun 14, 2007

TABLET; ORAL

LEXIVA

+ VIIV HLTHCARE

EQ 700MG BASE **

N021548 001 Oct 20, 2003

FOSAPREPITANT DIMEGLUMINE

POWDER; INTRAVENOUS

EMEND

+ MERCK AND CO INC

EQ 115MG BASE/VIAL **

N022023 001 Jan 25, 2008

FOSAPREPITANT DIMEGLUMINE

ACCORD HLTHCARE

EQ 150MG BASE/VIAL

A204025 001 Aug 26, 2020

APOTEX

EQ 150MG BASE/VIAL

A205020 001 Sep 05, 2019

ARTHUR GRP

EQ 150MG BASE/VIAL

A213199 001 Oct 04, 2021

GENEYORK PHARMS

EQ 150MG BASE/VIAL

A211624 001 Sep 05, 2019

MYLAN LABS LTD

EQ 115MG BASE/VIAL

A204015 001 Sep 05, 2019

NAVINTA LLC

EQ 150MG BASE/VIAL

A212957 002 Aug 20, 2020

SANDOZ

EQ 150MG BASE/VIAL

A203939 001 Dec 08, 2020

FOSCARNET SODIUM

INJECTABLE; INJECTION

FOSCARNET SODIUM

HOSPIRA

2.4GM/100ML

A077174 001 May 31, 2005

FOSFOMYCIN TROMETHAMINE

FOR SOLUTION; ORAL

MONUROL

+ ZAMBON SPA

EQ 3GM BASE/PACKET

N050717 001 Dec 19, 1996

FOSINOPRIL SODIUM

TABLET; ORAL

FOSINOPRIL SODIUM

ACTAVIS LABS FL INC

10MG

A076620 001 Oct 15, 2004

20MG

A076620 002 Oct 15, 2004

40MG

A076620 003 Oct 15, 2004

RANBAXY LABS LTD

10MG

A076580 001 Apr 23, 2004

20MG

A076580 002 Apr 23, 2004

40MG

A076580 003 Apr 23, 2004

UPSHER SMITH LABS

10MG

A076188 001 Oct 08, 2004

20MG

A076188 002 Oct 08, 2004

40MG

A076188 003 Oct 08, 2004

WATSON LABS

10MG

A076987 001 Dec 23, 2004

10MG

A077531 001 Aug 31, 2006

20MG

A076987 002 Dec 23, 2004

20MG

A077531 002 Aug 31, 2006

40MG

A076987 003 Dec 23, 2004

40MG

A077531 003 Aug 31, 2006

MONOPRIL

+ BRISTOL MYERS SQUIBB

10MG **

N019915 002 May 16, 1991

+

20MG **

N019915 003 May 16, 1991

+

40MG **

N019915 004 Mar 28, 1995

FOSINOPRIL SODIUM; HYDROCHLOROTHIAZIDE

TABLET; ORAL

FOSINOPRIL SODIUM AND HYDROCHLOROTHIAZIDE

ANI PHARMS

10MG;12.5MG

A076608 001 Dec 03, 2004

10MG;12.5MG

A077144 001 Aug 16, 2005

20MG;12.5MG

A076608 002 Dec 03, 2004

20MG;12.5MG

A077144 002 Aug 16, 2005

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

FOSINOPRIL SODIUM; HYDROCHLOROTHIAZIDE

TABLET; ORAL

FOSINOPRIL SODIUM AND HYDROCHLOROTHIAZIDE

AVET LIFESCIENCES	10MG;12.5MG	A079025 001	Sep 17, 2010
	20MG;12.5MG	A079025 002	Sep 17, 2010
MYLAN	10MG;12.5MG	A077705 001	Aug 14, 2006
	20MG;12.5MG	A077705 002	Aug 14, 2006
SUN PHARM INDS LTD	10MG;12.5MG	A076739 001	Dec 17, 2004
	20MG;12.5MG	A076739 002	Dec 17, 2004
TEVA	10MG;12.5MG	A076945 001	Jul 05, 2006
	20MG;12.5MG	A076945 002	Jul 05, 2006

MONOPRIL-HCT

+ BRISTOL MYERS SQUIBB	10MG;12.5MG **	N020286 002	Nov 30, 1994
+	20MG;12.5MG **	N020286 001	Nov 30, 1994

FOSNETUPITANT CHLORIDE HYDROCHLORIDE; PALONOSETRON HYDROCHLORIDE

POWDER; INTRAVENOUS

AKYNZEO

+ HELSINN HLTHCARE	EQ 235MG BASE/VIAL;EQ 0.25MG BASE/VIAL	N210493 001	Apr 19, 2018
--------------------	--	-------------	--------------

FOSPHENYTOIN SODIUM

INJECTABLE; INJECTION

FOSPHENYTOIN SODIUM

AM REGENT	EQ 50MG PHENYTOIN NA/ML	A078277 001	Aug 06, 2007
	EQ 50MG PHENYTOIN NA/ML	A090099 001	May 13, 2010
AMNEAL	EQ 50MG PHENYTOIN NA/ML	A078476 001	Mar 18, 2008
APOTEX INC	EQ 50MG PHENYTOIN NA/ML	A078126 001	Aug 06, 2007
DR REDDYS	EQ 50MG PHENYTOIN NA/ML	A076886 001	Aug 06, 2007
HOSPIRA	EQ 50MG PHENYTOIN NA/ML	A078158 001	Aug 06, 2007
NORVIUM BIOSCIENCE	EQ 50MG PHENYTOIN NA/ML	A078736 001	Jun 08, 2010

SOLUTION; INTRAVENOUS

SESQUIENT

+ LUPIN	EQ 100MG PHENYTOIN NA/2ML (EQ 50MG PHENYTOIN NA/ML)	N210864 001	Nov 05, 2020
+	EQ 500MG PHENYTOIN NA/10ML (EQ 50MG PHENYTOIN NA/ML)	N210864 002	Nov 05, 2020

FOSPROPOFOL DISODIUM

SOLUTION; INTRAVENOUS

LUSEDRA

EISAI INC	1050MG/30ML (35MG/ML)	N022244 001	Dec 12, 2008
-----------	-----------------------	-------------	--------------

FROVATRIPTAN SUCCINATE

TABLET; ORAL

FROVATRIPTAN SUCCINATE

NORVIUM BIOSCIENCE	EQ 2.5MG BASE	A202931 001	Aug 28, 2014
--------------------	---------------	-------------	--------------

FULVESTRANT

SOLUTION; INTRAMUSCULAR

FASLODEX

+ ASTRAZENECA	125MG/2.5ML (50MG/ML)	N021344 002	Apr 25, 2002
---------------	-----------------------	-------------	--------------

FULVESTRANT

APOTEX	250MG/5ML (50MG/ML)	A211730 001	Jun 11, 2021
AVYXA HOLDINGS	250MG/5ML (50MG/ML)	N210063 001	Aug 19, 2019

FURAZOLIDONE

SUSPENSION; ORAL

FUROXONE

SHIRE	50MG/15ML	N011323 002	
-------	-----------	-------------	--

TABLET; ORAL

FUROXONE

SHIRE	100MG	N011270 002	
-------	-------	-------------	--

FUROSEMIDE

INJECTABLE; INJECTION

FUROSEMIDE

ABRAXIS PHARM	10MG/ML	N018507 001	Jul 30, 1982
	10MG/ML	N019036 001	Aug 13, 1984
+ AM REGENT	10MG/ML **	N018579 001	Nov 30, 1983
ASTRAZENECA	10MG/ML	A070014 001	Sep 09, 1985
HIKMA	10MG/ML	A071439 001	Sep 14, 1990
HOSPIRA	10MG/ML	A070578 001	Jul 08, 1987
	10MG/ML	A072080 001	Aug 13, 1991

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

FUROSEMIDE

INJECTABLE; INJECTION

FUROSEMIDE

	10MG/ML	A074337 001	Oct 31, 1994
IGI LABS INC	10MG/ML	A070095 001	Sep 09, 1985
	10MG/ML	A070096 001	Sep 09, 1985
INTL MEDICATION	10MG/ML	N018025 001	
MARSAM PHARMS LLC	10MG/ML	A074017 001	Jun 30, 1994
SMITH AND NEPHEW	10MG/ML	A070023 001	Feb 05, 1986
	10MG/ML	A070078 001	Feb 05, 1986
WARNER CHILCOTT	10MG/ML	N018420 001	Feb 26, 1982
WATSON LABS	10MG/ML	A070019 001	Sep 22, 1986
	10MG/ML	A070604 001	Jan 02, 1987
WYETH AYERST	10MG/ML	N018670 001	Jul 20, 1982

LASIX

+ SANOFI AVENTIS US 10MG/ML **

N016363 001

SOLUTION; ORAL

FUROSEMIDE

PHARMOBEDIANT CNSLTG	10MG/ML	A070655 001	Oct 02, 1987
----------------------	---------	-------------	--------------

LASIX

SANOFI AVENTIS US 10MG/ML

N017688 001

TABLET; ORAL

FUROSEMIDE

ANI PHARMS	20MG	A071379 001	Jan 02, 1987
	40MG	A070413 001	Feb 26, 1986
	80MG	A071594 001	Feb 09, 1988
CHARTWELL RX	20MG	N018413 001	Nov 30, 1983
	40MG	N018413 002	Nov 30, 1983
EPIC PHARMA LLC	40MG	N018750 002	Jul 30, 1984
ESJAY PHARMA	20MG	N018415 001	Jul 27, 1982
	40MG	N018415 002	Jul 27, 1982
	80MG	N018415 003	Nov 26, 1984
INTL MEDICATION	20MG	N018753 001	Feb 28, 1984
	40MG	N018753 002	Feb 28, 1984
KALAPHARM	20MG	N018868 001	Jun 28, 1983
	40MG	N018868 002	Jun 28, 1983
MYLAN	20MG	N018487 001	
	40MG	N018487 002	
	80MG	A070082 001	Oct 29, 1986
SUN PHARM INDS INC	20MG	A091258 001	Apr 01, 2014
	40MG	A091258 002	Apr 01, 2014
	40MG	N018790 001	Nov 29, 1983
	80MG	A091258 003	Apr 01, 2014
SUN PHARM INDUSTRIES	20MG	A070043 001	Sep 26, 1985
	80MG	A070100 001	Jan 26, 1988
SUPERPHARM	20MG	N018370 002	Jun 26, 1984
	40MG	N018370 001	Feb 10, 1983
WARNER CHILCOTT	20MG	N018419 001	Jan 31, 1983
	40MG	N018419 002	Jan 31, 1983
	80MG	N018419 003	Nov 13, 1984
WATSON LABS	20MG	A070412 001	Feb 26, 1986
	20MG	N018369 001	May 14, 1982
	40MG	A070450 001	Nov 22, 1985
	40MG	N018369 002	May 14, 1982
WATSON LABS TEVA	20MG	A070449 001	Nov 22, 1985
	80MG	A070528 001	Jan 07, 1986

GABAPENTIN

CAPSULE; ORAL

GABAPENTIN

CHARTWELL RX	100MG	A091008 001	Oct 26, 2017
	300MG	A091008 002	Oct 26, 2017
	400MG	A091008 003	Oct 26, 2017
HIKMA	100MG	A078150 001	Sep 25, 2007
	300MG	A078150 002	Sep 25, 2007
	400MG	A078150 003	Sep 25, 2007
IPCA LABS LTD	100MG	A208928 001	Nov 20, 2023
	300MG	A208928 002	Nov 20, 2023
	400MG	A208928 003	Nov 20, 2023
NORVIUM BIOSCIENCE	100MG	A090158 001	Feb 14, 2011

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

GABAPENTIN

CAPSULE; ORAL

GABAPENTIN

	300MG	A090158	002	Feb 14, 2011
	400MG	A090158	003	Feb 14, 2011
SANDOZ	100MG	A075428	001	Jan 24, 2006
	100MG	A075539	001	Apr 06, 2005
	300MG	A075428	002	Jan 24, 2006
	300MG	A075539	002	Apr 06, 2005
	400MG	A075428	003	Jan 24, 2006
	400MG	A075539	003	Apr 06, 2005
SUN PHARM INDS LTD	100MG	A076606	001	Oct 07, 2005
	100MG	A077242	001	Aug 24, 2006
	300MG	A076606	002	Oct 07, 2005
	300MG	A077242	002	Aug 24, 2006
	400MG	A076606	003	Oct 07, 2005
	400MG	A077242	003	Aug 24, 2006
SUN PHARM INDUSTRIES	100MG	A076537	001	Jun 30, 2005
	300MG	A076537	002	Jun 30, 2005
	400MG	A076537	003	Jun 30, 2005
TEVA PHARMS	100MG	A075435	001	Oct 08, 2004
	300MG	A075435	002	Oct 08, 2004
	400MG	A075435	003	Oct 08, 2004
WATSON LABS	100MG	A075485	003	May 11, 2007
	300MG	A075485	002	May 11, 2007
	400MG	A075485	001	May 11, 2007

SOLUTION; ORAL

GABAPENTIN

PAI HOLDINGS PHARM	250MG/5ML	A211330	001	Dec 03, 2019
--------------------	-----------	---------	-----	--------------

TABLET; ORAL

GABAPENTIN

CHARTWELL RX	600MG	A076120	001	Jan 27, 2006
	800MG	A076120	002	Jan 27, 2006
HIKMA PHARMS	600MG	A078782	001	Jul 21, 2011
	800MG	A078782	002	Jul 21, 2011
INVAGEN PHARMS	600MG	A202764	001	Oct 16, 2012
	800MG	A202764	002	Oct 16, 2012
INVATECH	600MG	A076877	001	Jul 06, 2006
	800MG	A076877	002	Jul 06, 2006
IPCA LABS LTD	600MG	A209855	001	May 29, 2024
	800MG	A209855	002	May 29, 2024
IVAX SUB TEVA PHARMS	600MG	A076017	004	Apr 29, 2005
	800MG	A076017	005	Apr 29, 2005
LUPIN LTD	600MG	A209306	001	Aug 24, 2018
	800MG	A209306	002	Aug 24, 2018
NORVIUM BIOSCIENCE	600MG	A090335	001	Jun 01, 2010
	800MG	A090335	002	Jun 01, 2010
RANBAXY	600MG	A076605	001	Sep 14, 2005
	800MG	A076605	002	Sep 14, 2005
SUN PHARM INDS LTD	600MG	A077525	001	Aug 24, 2006
	800MG	A077525	002	Aug 24, 2006
TEVA	600MG	A075827	001	Dec 15, 2004
	800MG	A075827	002	Dec 15, 2004
TEVA PHARMS USA	600MG	A205807	001	Mar 10, 2017
	800MG	A205807	002	Mar 10, 2017

GADODIAMIDE

INJECTABLE; INJECTION

OMNISCAN

+ GE HEALTHCARE	14.35GM/50ML (287MG/ML)	N022066	001	Sep 05, 2007
-----------------	-------------------------	---------	-----	--------------

+	28.7GM/100ML (287MG/ML)	N022066	002	Sep 05, 2007
---	-------------------------	---------	-----	--------------

GADOFOSVESET TRISODIUM

SOLUTION; INTRAVENOUS

ABLAVAR

LANTHEUS MEDCL	2440MG/10ML (244MG/ML)	N021711	001	Dec 22, 2008
	3660MG/15ML (244MG/ML)	N021711	002	Dec 22, 2008

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

GADOPENTETATE DIMEGLUMINE

INJECTABLE; INJECTION

MAGNEVIST

+	BAYER HLTHCARE	469.01MG/ML	N019596 001	Jun 02, 1988
+		469.01MG/ML	N021037 001	Mar 10, 2000

GADOVERSETAMIDE

INJECTABLE; INJECTION

OPTIMARK

+	LIEBEL-FLARSHEIM	1654.5MG/5ML (330.9MG/ML)	N020937 001	Dec 08, 1999
+		3309MG/10ML (330.9MG/ML)	N020937 002	Dec 08, 1999
+		4963.5MG/15ML (330.9MG/ML)	N020937 003	Dec 08, 1999
+		6618MG/20ML (330.9MG/ML)	N020937 004	Dec 08, 1999
+		16.545GM/50ML (330.9MG/ML)	N020975 001	Dec 08, 1999

OPTIMARK IN PLASTIC CONTAINER

+	LIEBEL-FLARSHEIM	3309MG/10ML (330.9MG/ML)	N020976 002	Dec 08, 1999
+		4963.5MG/15ML (330.9MG/ML)	N020976 003	Dec 08, 1999
+		6618MG/20ML (330.9MG/ML)	N020976 004	Dec 08, 1999
+		9927MG/30ML (330.9MG/ML)	N020976 001	Dec 08, 1999

GALANTAMINE HYDROBROMIDE

CAPSULE, EXTENDED RELEASE; ORAL

GALANTAMINE HYDROBROMIDE

IMPAX LABS

		EQ 8MG BASE	A078484 001	May 27, 2009
		EQ 16MG BASE	A078484 002	May 27, 2009
		EQ 24MG BASE	A078484 003	May 27, 2009
	NORVIUM BIOSCIENCE	EQ 8MG BASE	A090900 001	Jan 24, 2011
		EQ 16MG BASE	A090900 002	Jan 24, 2011
		EQ 24MG BASE	A090900 003	Jan 24, 2011

RAZADYNE ER

+	JANSSEN PHARMS	EQ 8MG BASE **	N021615 001	Apr 01, 2005
+		EQ 16MG BASE **	N021615 002	Apr 01, 2005
+		EQ 24MG BASE **	N021615 003	Apr 01, 2005

SOLUTION; ORAL

RAZADYNE

	JANSSEN PHARMS	4MG/ML **	N021224 001	Jun 22, 2001
--	----------------	-----------	-------------	--------------

TABLET; ORAL

GALANTAMINE HYDROBROMIDE

APOTEX INC

		EQ 4MG BASE	A077781 001	Sep 27, 2011
		EQ 8MG BASE	A077781 002	Sep 27, 2011
		EQ 12MG BASE	A077781 003	Sep 27, 2011

CHARTWELL RX

		EQ 4MG BASE	A077587 001	Jul 09, 2009
		EQ 8MG BASE	A077587 002	Jul 09, 2009
		EQ 12MG BASE	A077587 003	Jul 09, 2009

HERITAGE PHARMA

		EQ 4MG BASE	A077585 001	Sep 15, 2009
		EQ 8MG BASE	A077585 002	Sep 15, 2009
		EQ 12MG BASE	A077585 003	Sep 15, 2009

HIKMA

		EQ 4MG BASE	A077608 001	Feb 11, 2009
		EQ 8MG BASE	A077608 002	Feb 11, 2009
		EQ 12MG BASE	A077608 003	Feb 11, 2009

MYLAN

		EQ 4MG BASE	A077590 001	May 29, 2009
		EQ 8MG BASE	A077590 002	May 29, 2009
		EQ 12MG BASE	A077590 003	May 29, 2009

NORVIUM BIOSCIENCE

		EQ 4MG BASE	A077603 001	Aug 28, 2008
		EQ 8MG BASE	A077603 002	Aug 28, 2008
		EQ 12MG BASE	A077603 003	Aug 28, 2008

RAZADYNE

+	JANSSEN PHARMS	EQ 4MG BASE **	N021169 001	Feb 28, 2001
+		EQ 8MG BASE **	N021169 002	Feb 28, 2001
+		EQ 12MG BASE **	N021169 003	Feb 28, 2001

GALLAMINE TRIETHIODIDE

INJECTABLE; INJECTION

FLAXEDIL

DAVIS AND GECK

		20MG/ML	N007842 001
		100MG/ML	N007842 002

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

GALLIUM CITRATE GA-67

INJECTABLE; INJECTION

GALLIUM CITRATE GA 67

GE HEALTHCARE 1mCi/ML N017700 001

LANTHEUS MEDCL 2mCi/ML N017478 001

NEOSCAN

GE HEALTHCARE 2mCi/ML N017655 001

GALLIUM NITRATE

INJECTABLE; INJECTION

GANITE

CHAPTER 7 TRUSTEE 25MG/ML ** N019961 002 Jan 17, 1991

GANCICLOVIR

CAPSULE; ORAL

CYTOVENE

+ ROCHE PALO 250MG ** N020460 001 Dec 22, 1994

+ 500MG ** N020460 002 Dec 12, 1997

GANCICLOVIR

RANBAXY LABS LTD 250MG A076457 001 Jun 27, 2003

500MG A076457 002 Jun 27, 2003

IMPLANT; IMPLANTATION

VITRASERT

BAUSCH AND LOMB 4.5MG N020569 001 Mar 04, 1996

GANCICLOVIR SODIUM

INJECTABLE; INJECTION

CYTOVENE

+ CHEPLAPHARM EQ 500MG BASE/VIAL ** N019661 001 Jun 23, 1989

GANCICLOVIR SODIUM

AM REGENT EQ 500MG BASE/VIAL A202624 001 Sep 18, 2013

CUSTOPHARM INC EQ 500MG BASE/VIAL A212001 001 Jun 20, 2019

STERISCIENCE SPECLTS EQ 500MG BASE/VIAL A204560 001 Nov 17, 2017

GATIFLOXACIN

SOLUTION/DROPS; OPHTHALMIC

GATIFLOXACIN

APOTEX INC 0.3% A079084 001 Aug 19, 2011

HIKMA 0.5% A203189 001 Sep 03, 2014

RISING 0.5% A206446 001 Jun 08, 2018

TORRENT 0.5% A213542 001 Nov 03, 2021

ZYMAR

+ ALLERGAN 0.3% N021493 001 Mar 28, 2003

GEFITINIB

TABLET; ORAL

IRESSA

ASTRAZENECA 250MG N021399 001 May 05, 2003

GEMCITABINE HYDROCHLORIDE

INJECTABLE; INJECTION

GEMCITABINE HYDROCHLORIDE

ACCORD HLTHCARE EQ 200MG BASE/VIAL A091594 001 Jul 25, 2011

EQ 1GM BASE/VIAL A091594 002 Jul 25, 2011

EQ 2GM BASE/VIAL A091594 003 Jul 25, 2011

ACTAVIS INC 200MG/5.26ML (38MG/ML) A204549 001 Apr 11, 2016

1GM/26.3ML (38MG/ML) A204549 002 Apr 11, 2016

2GM/52.6ML (38MG/ML) A204549 003 Apr 11, 2016

ACTAVIS TOTOWA EQ 200MG BASE/VIAL A079160 001 Jul 25, 2011

EQ 1GM BASE/VIAL A079160 002 Jul 25, 2011

EQ 2GM BASE/VIAL A079160 003 Jul 28, 2016

AM REGENT EQ 200MG BASE/VIAL A202031 001 May 07, 2013

EQ 1GM BASE/VIAL A202031 002 May 07, 2013

APOTEX 200MG/5.26ML (38MG/ML) A206776 001 May 23, 2017

1GM/26.3ML (38MG/ML) A206776 002 May 23, 2017

2GM/52.6ML (38MG/ML) A206776 003 May 23, 2017

EMCURE PHARMS LTD EQ 200MG BASE/VIAL A202063 001 Sep 11, 2012

EQ 1GM BASE/VIAL A202063 002 Sep 11, 2012

HAMELN RDS GMBH EQ 200MG BASE/VIAL A090663 001 Sep 10, 2012

EQ 1GM BASE/VIAL A090663 002 Sep 10, 2012

NORVIUM BIOSCIENCE EQ 200MG BASE/VIAL A200145 001 Jul 25, 2011

EQ 1GM BASE/VIAL A200145 002 Jul 25, 2011

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

GEMCITABINE HYDROCHLORIDE

INJECTABLE; INJECTION

GEMCITABINE HYDROCHLORIDE

	EQ 2GM BASE/VIAL	A200145 003	Jul 25, 2011
SAGENT PHARMS	EQ 200MG BASE/VIAL	A091597 001	May 07, 2013
	EQ 1GM BASE/VIAL	A091597 002	May 07, 2013
TEVA PHARMS	EQ 200MG BASE/VIAL	A077983 002	Jan 25, 2011
	EQ 1GM BASE/VIAL	A077983 001	Jan 25, 2011
GEMZAR			
+ LILLY	EQ 200MG BASE/VIAL **	N020509 001	May 15, 1996
+	EQ 1GM BASE/VIAL **	N020509 002	May 15, 1996

SOLUTION; INTRAVENOUS

INFUGEM

+ SUN PHARM	EQ 1200MG BASE/120ML (EQ 10MG BASE/ML)	N208313 001	Jul 16, 2018
+	EQ 1300MG BASE/130ML (EQ 10MG BASE/ML)	N208313 002	Jul 16, 2018
+	EQ 1400MG BASE/140ML (EQ 10MG BASE/ML)	N208313 003	Jul 16, 2018
+	EQ 1500MG BASE/150ML (EQ 10MG BASE/ML)	N208313 004	Jul 16, 2018
+	EQ 1600MG BASE/160ML (EQ 10MG BASE/ML)	N208313 005	Jul 16, 2018
+	EQ 1700MG BASE/170ML (EQ 10MG BASE/ML)	N208313 006	Jul 16, 2018
+	EQ 1800MG BASE/180ML (EQ 10MG BASE/ML)	N208313 007	Jul 16, 2018
+	EQ 1900MG BASE/190ML (EQ 10MG BASE/ML)	N208313 008	Jul 16, 2018
+	EQ 2000MG BASE/200ML (EQ 10MG BASE/ML)	N208313 009	Jul 16, 2018
+	EQ 2200MG BASE/220ML (EQ 10MG BASE/ML)	N208313 010	Jul 16, 2018

GEMFIBROZIL

CAPSULE; ORAL

GEMFIBROZIL

NATCO	300MG	A073466 001	Jan 25, 1993
PUREPAC PHARM	300MG	A072929 001	Jan 29, 1993
LOPID			
PFIZER PHARMS	200MG	N018422 001	
	300MG	N018422 002	

TABLET; ORAL

GEMFIBROZIL

CADILA	600MG	A204189 001	Aug 28, 2018
CHARTWELL RX	600MG	A074615 001	Sep 29, 1995
HIKMA PHARMS	600MG	A078599 001	Aug 16, 2010
NORVIUM BIOSCIENCE	600MG	A074452 001	Feb 16, 1995
PUREPAC PHARM	600MG	A074360 001	Aug 31, 1994
SUN PHARM INDS INC	600MG	A079239 001	Dec 29, 2008
TEVA	600MG	A074256 001	Oct 31, 1993
WATSON LABS	600MG	A074156 001	Oct 24, 1994
	600MG	A074442 001	Apr 28, 1995

GEMIFLOXACIN MESYLATE

TABLET; ORAL

FACTIVE

+ LG CHEM LTD	EQ 320MG BASE **	N021158 001	Apr 04, 2003
GEMIFLOXACIN MESYLATE			
ORBION PHARMS	EQ 320MG BASE	A090466 001	Jun 15, 2015

GENTAMICIN SULFATE

CREAM; TOPICAL

GARAMYCIN

SCHERING	EQ 0.1% BASE **	A060462 001	
GENTAFAIR			
PHARMAFAIR	EQ 0.1% BASE	A062458 001	Sep 01, 1983
GENTAMICIN SULFATE			
ALPHARMA US PHARMS	EQ 0.1% BASE	A062471 001	Sep 27, 1983
FOUGERA PHARMS INC	EQ 0.1% BASE	A062531 001	Jul 05, 1984
PHARMADERM	EQ 1MG BASE/GM	A062530 001	Jul 05, 1984
TARO	EQ 0.1% BASE	A062427 001	May 26, 1983
INJECTABLE; INJECTION			
APOGEN			
KING PHARMS	EQ 10MG BASE/ML	A062289 001	
	EQ 40MG BASE/ML	A062289 002	
BRISTAGEN			
BRISTOL	EQ 40MG BASE/ML	A062288 001	
GARAMYCIN			
SCHERING	EQ 1MG BASE/ML **	A061716 002	
	EQ 10MG BASE/ML **	A061739 001	

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

GENTAMICIN SULFATE

INJECTABLE; INJECTION

GARAMYCIN

EQ 40MG BASE/ML ** A061716 001

GENTAFAIR

PHARMAFAIR

EQ 40MG BASE/ML A062493 001 Aug 28, 1985

GENTAMICIN

INTL MEDICATION

EQ 1MG BASE/ML A062325 003 Jun 23, 1982

EQ 40MG BASE/ML A062325 001

EQ 100MG BASE/100ML A062325 004 Jun 23, 1982

GENTAMICIN SULFATE

ABBOTT

EQ 1.2MG BASE/ML A062413 001 Aug 11, 1983

EQ 1.4MG BASE/ML A062413 002 Aug 11, 1983

EQ 1.6MG BASE/ML A062413 003 Aug 11, 1983

EQ 1.8MG BASE/ML A062413 004 Aug 11, 1983

EQ 2MG BASE/ML A062413 005 Aug 11, 1983

EQ 60MG BASE/100ML A062413 006 Aug 11, 1983

EQ 70MG BASE/100ML A062413 007 Aug 11, 1983

EQ 80MG BASE/100ML A062413 008 Aug 11, 1983

EQ 90MG BASE/100ML A062413 009 Aug 11, 1983

EQ 100MG BASE/100ML A062413 010 Aug 11, 1983

FRESENIUS KABI USA

EQ 10MG BASE/ML A062356 001 Mar 04, 1982

EQ 40MG BASE/ML A062356 002 Mar 04, 1982

HOSPIRA

EQ 10MG BASE/ML A062612 004 Feb 20, 1986

KALAPHARM

EQ 40MG BASE/ML A062354 001 Apr 05, 1982

PHARM SPEC

EQ 40MG BASE/ML A062340 001 Mar 28, 1983

SOLOPAK

EQ 10MG BASE/ML A062507 001 Jun 06, 1985

EQ 40MG BASE/ML A062507 002 Jun 06, 1985

TEVA PARENTERAL

EQ 10MG BASE/ML A063149 001 Nov 21, 1991

EQ 40MG BASE/ML A063106 002 Nov 21, 1991

WATSON LABS

EQ 10MG BASE/ML A062318 002

EQ 40MG BASE/ML A062318 001

WYETH AYERST

EQ 10MG BASE/ML A062264 001

EQ 40MG BASE/ML A062264 002

GENTAMICIN SULFATE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

B BRAUN

EQ 0.8MG BASE/ML A062814 001 Aug 28, 1987

EQ 1.2MG BASE/ML A062814 002 Aug 28, 1987

EQ 1.4MG BASE/ML A062814 003 Aug 28, 1987

EQ 1.6MG BASE/ML A062814 004 Aug 28, 1987

EQ 1.8MG BASE/ML A062814 005 Aug 28, 1987

EQ 2MG BASE/ML A062814 006 Aug 28, 1987

EQ 2.4MG BASE/ML A062814 007 Aug 28, 1987

EQ 40MG BASE/100ML A062814 008 Aug 28, 1987

EQ 60MG BASE/100ML A062814 009 Aug 28, 1987

EQ 70MG BASE/100ML A062814 010 Aug 28, 1987

EQ 80MG BASE/100ML A062814 011 Aug 28, 1987

EQ 90MG BASE/100ML A062814 012 Aug 28, 1987

EQ 100MG BASE/100ML A062814 013 Aug 28, 1987

EQ 120MG BASE/100ML A062814 014 Aug 28, 1987

BAXTER HLTHCARE

EQ 0.8MG BASE/ML A062373 001 Sep 07, 1982

EQ 2.4MG BASE/ML A062373 010 Sep 07, 1982

EQ 40MG BASE/100ML A062373 003 Sep 07, 1982

EQ 60MG BASE/100ML A062373 004 Sep 07, 1982

HOSPIRA

EQ 1.2MG BASE/ML A062588 001 Jan 06, 1986

EQ 1.4MG BASE/ML A062414 002 Aug 15, 1983

EQ 1.4MG BASE/ML A062588 002 Jan 06, 1986

EQ 1.6MG BASE/ML A062588 003 Jan 06, 1986

EQ 1.8MG BASE/ML A062414 004 Aug 15, 1983

EQ 1.8MG BASE/ML A062588 004 Jan 06, 1986

EQ 2MG BASE/ML A062414 005 Aug 15, 1983

EQ 2MG BASE/ML A062588 005 Jan 06, 1986

EQ 60MG BASE/100ML A062414 006 Aug 15, 1983

EQ 60MG BASE/100ML A062588 006 Jan 06, 1986

EQ 70MG BASE/100ML A062414 007 Aug 15, 1983

EQ 70MG BASE/100ML A062588 007 Jan 06, 1986

EQ 80MG BASE/100ML A062588 008 Jan 06, 1986

EQ 90MG BASE/100ML A062414 009 Aug 15, 1983

EQ 90MG BASE/100ML A062588 009 Jan 06, 1986

EQ 100MG BASE/100ML A062588 010 Jan 06, 1986

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

GENTAMICIN SULFATE

INJECTABLE; INJECTION

U-GENCIN

PHARMACIA AND UPJOHN	EQ 10MG BASE/ML	A062248	001	
	EQ 40MG BASE/ML	A062248	002	

INJECTABLE; INTRATHECAL

GARAMYCIN

+ SCHERING	EQ 2MG BASE/ML **	N050505	001	
------------	-------------------	---------	-----	--

OINTMENT; OPHTHALMIC

GARAMYCIN

SCHERING	EQ 0.3% BASE	N050425	001	
----------	--------------	---------	-----	--

GENTACIDIN

NOVARTIS	EQ 0.3% BASE	A062501	001	Jul 26, 1984
----------	--------------	---------	-----	--------------

GENTAFAIR

PHARMAFAIR	EQ 3MG BASE/GM	A062443	001	May 26, 1983
------------	----------------	---------	-----	--------------

GENTAMICIN SULFATE

SCIEGEN PHARMS INC	EQ 0.3% BASE	A064093	001	Aug 31, 1995
--------------------	--------------	---------	-----	--------------

OINTMENT; TOPICAL

GARAMYCIN

SCHERING	EQ 0.1% BASE **	A060463	001	
----------	-----------------	---------	-----	--

GENTAFAIR

PHARMAFAIR	EQ 0.1% BASE	A062444	001	May 26, 1983
------------	--------------	---------	-----	--------------

GENTAMICIN SULFATE

ALPHARMA US PHARMS	EQ 0.1% BASE	A062496	001	Mar 14, 1984
--------------------	--------------	---------	-----	--------------

ENCUBE	EQ 0.1% BASE	A209233	001	Dec 31, 2018
--------	--------------	---------	-----	--------------

PHARMADERM	EQ 0.1% BASE	A062534	001	Oct 10, 1984
------------	--------------	---------	-----	--------------

SOLUTION/DROPS; OPHTHALMIC

GARAMYCIN

+ SCHERING	EQ 0.3% BASE **	N050039	002	
------------	-----------------	---------	-----	--

GENOPTIC

ALLERGAN	EQ 0.3% BASE	A062452	001	Oct 10, 1984
----------	--------------	---------	-----	--------------

GENTACIDIN

NOVARTIS	EQ 0.3% BASE	A062480	001	Mar 30, 1984
----------	--------------	---------	-----	--------------

GENTAFAIR

PHARMAFAIR	EQ 0.3% BASE	A062440	001	May 03, 1983
------------	--------------	---------	-----	--------------

GENTAK

RENOVA PHARMS	EQ 0.3% BASE	A064163	001	Oct 12, 2001
---------------	--------------	---------	-----	--------------

GENTAMICIN SULFATE

ALCON PHARMS LTD	EQ 0.3% BASE	A062523	001	Nov 25, 1985
------------------	--------------	---------	-----	--------------

EPIC PHARMA LLC	EQ 0.3% BASE	A062635	001	Jan 08, 1987
-----------------	--------------	---------	-----	--------------

PACO	EQ 3MG BASE/ML	A062932	001	Nov 07, 1988
------	----------------	---------	-----	--------------

GENTAMICIN SULFATE; PREDNISOLONE ACETATE

OINTMENT; OPHTHALMIC

PRED-G

+ ALLERGAN	EQ 0.3% BASE; 0.6%	N050612	001	Dec 01, 1989
------------	--------------------	---------	-----	--------------

SUSPENSION/DROPS; OPHTHALMIC

PRED-G

+ ALLERGAN	EQ 0.3% BASE; 1%	N050586	001	Jun 10, 1988
------------	------------------	---------	-----	--------------

GENTIAN VIOLET

SUPPOSITORY; VAGINAL

GVS

SAVAGE LABS	0.4%	A083513	001	
-------------	------	---------	-----	--

TAMPON; VAGINAL

GENAPAX

KEY PHARMS	5MG	A085017	001	
------------	-----	---------	-----	--

GEPİRONE HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL

EXXUA

+ FABRE KRAMER	EQ 18.2MG BASE	N021164	001	Sep 22, 2023
----------------	----------------	---------	-----	--------------

	EQ 36.3MG BASE	N021164	002	Sep 22, 2023
--	----------------	---------	-----	--------------

	EQ 54.5MG BASE	N021164	003	Sep 22, 2023
--	----------------	---------	-----	--------------

	EQ 72.6MG BASE	N021164	004	Sep 22, 2023
--	----------------	---------	-----	--------------

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

GLATIRAMER ACETATEFOR SOLUTION;SUBCUTANEOUS
COPAXONE

+ TEVA PHARMS USA 20MG/VIAL

N020622 001 Dec 20, 1996

GLIMEPIRIDE

TABLET;ORAL

GLIMEPIRIDE

ACTAVIS LABS FL INC 1MG

A076995 001 Apr 27, 2010

2MG

A076995 002 Apr 27, 2010

4MG

A076995 003 Apr 27, 2010

EPIC PHARMA LLC 1MG

A077274 001 Oct 06, 2005

2MG

A077274 002 Oct 06, 2005

4MG

A077274 003 Oct 06, 2005

HIKMA PHARMS 1MG

A078952 001 Aug 01, 2013

2MG

A078952 002 Aug 01, 2013

4MG

A078952 003 Aug 01, 2013

MYLAN 1MG

A077486 001 Feb 10, 2006

2MG

A077486 002 Feb 10, 2006

4MG

A077486 003 Feb 10, 2006

NORVIUM BIOSCIENCE 1MG

A077624 001 Nov 28, 2005

2MG

A077624 002 Nov 28, 2005

4MG

A077624 003 Nov 28, 2005

RANBAXY 3MG

A077366 001 Oct 06, 2005

6MG

A077366 002 Oct 06, 2005

RANBAXY LABS LTD 1MG

A076875 001 Oct 06, 2005

2MG

A076875 002 Oct 06, 2005

4MG

A076875 003 Oct 06, 2005

8MG

A076875 004 Oct 06, 2005

TEVA 1MG

A076802 001 Oct 06, 2005

2MG

A076802 002 Oct 06, 2005

4MG

A076802 003 Oct 06, 2005

WATSON LABS 1MG

A077280 001 Feb 03, 2006

2MG

A077280 002 Feb 03, 2006

4MG

A077280 003 Feb 03, 2006

GLIMEPIRIDE; ROSIGLITAZONE MALEATE

TABLET;ORAL

AVANDARYL

+ SB PHARMCO 1MG;4MG **

N021700 001 Nov 23, 2005

+ 2MG;4MG **

N021700 002 Nov 23, 2005

+ 2MG;8MG **

N021700 004 Mar 30, 2007

+ 4MG;4MG **

N021700 003 Nov 23, 2005

+ 4MG;8MG **

N021700 005 Mar 30, 2007

ROSIGLITAZONE MALEATE AND GLIMEPIRIDE

TEVA PHARMS USA 1MG;4MG

A078709 001 Apr 01, 2016

2MG;4MG

A078709 002 Apr 01, 2016

2MG;8MG

A078709 004 Apr 01, 2016

4MG;4MG

A078709 003 Apr 01, 2016

4MG;8MG

A078709 005 Apr 01, 2016

GLIPIZIDE

TABLET;ORAL

GLIPIZIDE

ACCORD HLTHCARE 5MG

A074550 001 Sep 11, 1997

10MG

A074550 002 Sep 11, 1997

ANI PHARMS 5MG

A074387 001 Mar 04, 1996

10MG

A074387 002 Mar 04, 1996

BARR LABS INC 5MG

A074619 001 Apr 04, 1997

10MG

A074619 002 Apr 04, 1997

CHARTWELL RX 5MG

A074542 001 Jun 20, 1995

10MG

A074542 002 Jun 20, 1995

NORVIUM BIOSCIENCE 5MG

A074438 001 Jun 20, 1995

10MG

A074438 002 Jun 20, 1995

OXFORD PHARMS 5MG

A074378 001 Nov 28, 1994

10MG

A074378 002 Nov 28, 1994

SANDOZ 5MG

A074305 001 Apr 07, 1995

10MG

A074305 002 Apr 07, 1995

SUN PHARM INDS INC 5MG

A077820 001 Jul 11, 2006

10MG

A077820 002 Jul 11, 2006

WATSON LABS 5MG

A074370 001 Nov 22, 1994

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

GLIPIZIDE

TABLET; ORAL

GLIPIZIDE

10MG

A074370 002 Nov 22, 1994

GLUCOTROL

+ PFIZER

2.5MG **

N017783 003 May 11, 1993

+

5MG **

N017783 001 May 08, 1984

+

10MG **

N017783 002 May 08, 1984

TABLET, EXTENDED RELEASE; ORAL

GLIPIZIDE

ENDO OPERATIONS

5MG

A076159 002 Sep 20, 2013

10MG

A076159 001 Sep 20, 2013

NORVIUM BIOSCIENCE

2.5MG

A202298 001 May 19, 2015

5MG

A202298 002 May 19, 2015

10MG

A202298 003 May 19, 2015

GLUCOTROL XL

+ PFIZER

5MG

N020329 001 Apr 26, 1994

GLIPIZIDE; METFORMIN HYDROCHLORIDE

TABLET; ORAL

GLIPIZIDE AND METFORMIN HYDROCHLORIDE

ESJAY PHARMA

2.5MG; 250MG

A078083 001 Apr 12, 2007

2.5MG; 500MG

A078083 002 Apr 12, 2007

5MG; 500MG

A078083 003 Apr 12, 2007

SUN PHARM INDS INC

2.5MG; 250MG

A077620 001 Jan 11, 2008

2.5MG; 500MG

A077620 002 Jan 11, 2008

5MG; 500MG

A077620 003 Jan 11, 2008

METAGLIP

+ BRISTOL MYERS SQUIBB

2.5MG; 250MG **

N021460 001 Oct 21, 2002

+

2.5MG; 500MG **

N021460 002 Oct 21, 2002

+

5MG; 500MG **

N021460 003 Oct 21, 2002

GLUCAGON

INJECTABLE; INJECTION

GLUCAGON

+ LILLY

1MG/VIAL **

N020928 001 Sep 11, 1998

SOLUTION; SUBCUTANEOUS

GVOKE PFS

+ XERIS

0.5MG/0.1ML (0.5MG/0.1ML)

N212097 001 Sep 10, 2019

GLUCAGON HYDROCHLORIDE

INJECTABLE; INJECTION

GLUCAGON

+ LILLY

EQ 1MG BASE/VIAL **

N012122 001

+

EQ 10MG BASE/VIAL **

N012122 002

POWDER; INTRAMUSCULAR, INTRAVENOUS

GLUCAGEN

+ NOVO NORDISK

EQ 1MG BASE/VIAL **

N020918 001 Jun 22, 1998

POWDER; INTRAMUSCULAR, INTRAVENOUS, SUBCUTANEOUS

GLUCAGEN

+ NOVO NORDISK

EQ 1MG BASE/VIAL **

N020918 002 Jun 22, 1998

GLUTETHIMIDE

CAPSULE; ORAL

DORIDEN

SANOFI AVENTIS US

500MG

N009519 008

TABLET; ORAL

DORIDEN

SANOFI AVENTIS US

250MG

N009519 002

500MG

N009519 005

GLUTETHIMIDE

HALSEY

250MG

A089458 001 Oct 10, 1986

500MG

A089459 001 Oct 10, 1986

LANNETT

250MG

A083475 001

500MG

A085571 001

UCB INC

500MG

A085171 001

UPSHER SMITH LABS

500MG

A083234 002

VITARINE

500MG

A087297 001

WATSON LABS

500MG

A084362 001

500MG

A085763 001

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

GLYBURIDE

TABLET; ORAL

GLYBURIDE

ACTAVIS ELIZABETH	1.5MG	A075947 001	Nov 14, 2002
	3MG	A075947 002	Nov 14, 2002
	6MG	A075947 003	Nov 14, 2002
AUROBINDO PHARMA	1.25MG	A077537 001	Oct 18, 2007
	2.5MG	A077537 002	Oct 18, 2007
	5MG	A077537 003	Oct 18, 2007
CHARTWELL RX	1.25MG	A203581 001	Apr 14, 2016
	2.5MG	A203581 002	Apr 14, 2016
	5MG	A203581 003	Apr 14, 2016

GLYBURIDE (MICRONIZED)

CHARTWELL RX	1.5MG	A075174 001	Jun 22, 1998
	3MG	A075174 002	Jun 22, 1998
HIKMA	1.5MG	A075890 001	Jul 31, 2003
	3MG	A075890 002	Jul 31, 2003
	6MG	A075890 003	Jul 31, 2003
NATCO PHARMA	1.5MG	A074792 001	Jun 26, 1998
	3MG	A074792 002	Jun 26, 1998
	6MG	A074792 003	Aug 17, 1999
SANOFI AVENTIS US	1.5MG	N020055 001	Apr 17, 1992
	3MG	N020055 002	Apr 17, 1992
	6MG	N020055 003	Mar 08, 2000
STRIDES PHARMA	1.5MG	A074591 001	Dec 22, 1997
	3MG	A074591 002	Dec 22, 1997
	4.5MG	A074591 003	Dec 22, 1997
	6MG	A074591 004	Dec 22, 1997

GLYNASE

+ PFIZER 4.5MG ** N020051 003 Sep 24, 1993

MICRONASE

+ PFIZER 1.25MG ** N017498 001 May 01, 1984

+ PFIZER 2.5MG N017498 002 May 01, 1984

+ PFIZER 5MG ** N017498 003 May 01, 1984

GLYBURIDE; METFORMIN HYDROCHLORIDE

TABLET; ORAL

GLUCOVANCE

+ BRISTOL MYERS SQUIBB 1.25MG; 250MG ** N021178 001 Jul 31, 2000

+ BRISTOL MYERS SQUIBB 2.5MG; 500MG ** N021178 002 Jul 31, 2000

+ BRISTOL MYERS SQUIBB 5MG; 500MG ** N021178 003 Jul 31, 2000

GLYBURIDE AND METFORMIN HYDROCHLORIDE

IMPAX LABS INC	1.25MG; 250MG	A076731 001	Nov 19, 2004
	2.5MG; 500MG	A076731 002	Nov 19, 2004
	5MG; 500MG	A076731 003	Nov 19, 2004
TEVA	1.25MG; 250MG	A076821 001	Jan 27, 2005
	2.5MG; 500MG	A076821 002	Jan 27, 2005
	5MG; 500MG	A076821 003	Jan 27, 2005

GLYCEROL PHENYLBUTYRATE

LIQUID; ORAL

GLYCEROL PHENYLBUTYRATE

ENDO OPERATIONS 1.1GM/ML A205742 001 Dec 02, 2021

GLYCINE

SOLUTION; IRRIGATION

GLYCINE 1.5% IN PLASTIC CONTAINER

BAXTER HLTHCARE 1.5GM/100ML N018522 001 Feb 19, 1982

HOSPIRA 1.5GM/100ML N017633 001

GLYCOPYRROLATE

INJECTABLE; INJECTION

GLYCOPYRROLATE

ABRAXIS PHARM	0.2MG/ML	A088475 001	Jun 12, 1984
EUGIA PHARMA	0.2MG/ML	A210244 001	Nov 28, 2018
HOSPIRA	0.2MG/ML	A089393 001	Jun 15, 1988
LUPIN LTD	0.2MG/ML	A213655 001	Feb 07, 2023
MANKIND PHARMA	0.2MG/ML	A217354 001	Feb 27, 2023
TEVA PARENTERAL	0.2MG/ML	A081169 001	Sep 10, 1991
WATSON LABS	0.2MG/ML	A086947 001	Jun 24, 1983
ZYDUS PHARMS	0.2MG/ML	A214213 001	Nov 09, 2021

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

GLYCOPYRROLATE

INJECTABLE; INJECTION

ROBINUL

+	HIKMA	0.2MG/ML **	N017558	001	
	ROBINS AH	0.2MG/ML	N014764	001	

POWDER; INHALATION

SEEBRI

+	NOVARTIS	15.6MCG/INH	N207923	001	Oct 29, 2015
---	----------	-------------	---------	-----	--------------

SOLUTION; INHALATION

LONHALA MAGNAIR KIT

+	SUMITOMO PHARMA AM	25MCG/ML	N208437	001	Dec 05, 2017
---	--------------------	----------	---------	-----	--------------

SOLUTION; INTRAMUSCULAR, INTRAVENOUS

GLYRX-PF

+	EXELA PHARMA	0.6MG/3ML (0.2MG/ML)	N210997	004	Dec 14, 2020
---	--------------	----------------------	---------	-----	--------------

TABLET; ORAL

GLYCOPYRROLATE

CHARTWELL RX

1MG	A040568	001	Dec 22, 2004
2MG	A040568	002	Dec 22, 2004
1MG	A040836	001	Mar 05, 2009
2MG	A040836	002	Mar 05, 2009
1MG	A085562	001	
1MG	A086902	001	
2MG	A085563	001	
2MG	A086178	001	
2MG	A086900	001	

TABLET, ORALLY DISINTEGRATING; ORAL

DARTISLA ODT

EDENBRIDGE PHARMS	0.85MG	N215019	002	Oct 11, 2023
-------------------	--------	---------	-----	--------------

+		1.7MG	N215019	001	Dec 16, 2021
---	--	-------	---------	-----	--------------

GLYCOPYRROLATE; INDACATEROL MALEATE

POWDER; INHALATION

UTIBRON

+	NOVARTIS	15.6MCG/INH; 27.5MCG/INH	N207930	001	Oct 29, 2015
---	----------	--------------------------	---------	-----	--------------

GONADORELIN ACETATE

INJECTABLE; INJECTION

LUTREPULSE KIT

FERRING	0.8MG/VIAL	N019687	001	Oct 10, 1989
	3.2MG/VIAL	N019687	002	Oct 10, 1989

GONADORELIN HYDROCHLORIDE

INJECTABLE; INJECTION

FACTREL

HIKMA	EQ 0.1MG BASE/VIAL	N018123	001	Sep 30, 1982
	EQ 0.2MG BASE/VIAL	N018123	002	Sep 30, 1982
	EQ 0.5MG BASE/VIAL	N018123	003	Sep 30, 1982

GRAMICIDIN; NEOMYCIN SULFATE; POLYMYXIN B SULFATE

SOLUTION/DROPS; OPHTHALMIC

NEO-POLYCIN

DOW PHARM	0.025MG/ML; EQ 1.75MG BASE/ML; 10,000 UNITS/ML	A060427	001	
-----------	--	---------	-----	--

NEOMYCIN AND POLYMYXIN B SULFATES AND GRAMICIDIN

IPHARM	0.025MG/ML; EQ 1.75MG BASE/ML; 10,000 UNITS/ML	A062818	001	Oct 11, 1988
--------	--	---------	-----	--------------

WATSON LABS	0.025MG/ML; EQ 1.75MG BASE/ML; 10,000 UNITS/ML	A062788	001	Jun 11, 1987
-------------	--	---------	-----	--------------

NEOMYCIN SULFATE AND POLYMYXIN B SULFATE GRAMICIDIN

PHARMAFAIR	0.025MG/ML; EQ 1.75MG BASE/ML; 10,000 UNITS/ML	A062383	001	Aug 31, 1982
------------	--	---------	-----	--------------

GRANISETRON HYDROCHLORIDE

INJECTABLE; INJECTION

GRANISETRON HYDROCHLORIDE

AM REGENT	EQ 1MG BASE/ML (EQ 1MG BASE/ML)	A091274	001	Sep 22, 2010
BAXTER HLTHCARE CORP	EQ 0.1MG BASE/ML (EQ 0.1MG BASE/ML)	A078197	001	Dec 31, 2007
	EQ 1MG BASE/ML (EQ 1MG BASE/ML)	A078198	001	Jun 30, 2008
	EQ 4MG BASE/4ML (EQ 1MG BASE/ML)	A078198	002	Jun 30, 2008
EPIC PHARMA LLC	EQ 0.1MG BASE/ML (EQ 0.1MG BASE/ML)	A079119	001	Sep 10, 2009
EUGIA PHARMA	EQ 1MG BASE/ML (EQ 1MG BASE/ML)	A204238	001	Jul 06, 2016
	EQ 4MG BASE/4ML (EQ 1MG BASE/ML)	A204238	002	Jul 06, 2016

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

GRANISETRON HYDROCHLORIDE

INJECTABLE; INJECTION

GRANISETRON HYDROCHLORIDE

NORVIUM BIOSCIENCE	EQ 0.1MG BASE/ML (EQ 0.1MG BASE/ML)	A203454 001	Apr 04, 2017
	EQ 1MG BASE/ML (EQ 1MG BASE/ML)	A203454 002	Apr 04, 2017
	EQ 4MG BASE/4ML (EQ 1MG BASE/ML)	A203453 001	Jan 31, 2017
SANDOZ	EQ 0.1MG BASE/ML (EQ 0.1MG BASE/ML)	A078808 001	Apr 29, 2008
TEVA PHARMS USA	EQ 1MG BASE/ML (EQ 1MG BASE/ML)	A077963 001	Jan 03, 2008
WOCKHARDT USA	EQ 0.1MG BASE/ML (EQ 0.1MG BASE/ML)	A078566 001	Feb 29, 2008
	EQ 1MG BASE/ML (EQ 1MG BASE/ML)	A078564 001	Jun 30, 2008
	EQ 4MG BASE/4ML (EQ 1MG BASE/ML)	A078565 001	Jun 30, 2008
YUNG SHIN PHARM	EQ 0.1MG BASE/ML (EQ 0.1MG BASE/ML)	A202647 001	Mar 06, 2020
	EQ 1MG BASE/ML (EQ 1MG BASE/ML)	A202648 001	Jun 29, 2020
	EQ 4MG BASE/4ML (EQ 1MG BASE/ML)	A202648 002	Jun 29, 2020

GRANISETRON HYDROCHLORIDE PRESERVATIVE FREE

DR REDDYS	EQ 1MG BASE/ML (EQ 1MG BASE/ML)	A077165 001	Dec 31, 2007
-----------	---------------------------------	-------------	--------------

KYTRIL

+ ROCHE	EQ 0.1MG BASE/ML (EQ 0.1MG BASE/ML) **	N020239 003	Sep 17, 2004
+	EQ 1MG BASE/ML (EQ 1MG BASE/ML) **	N020239 004	Mar 11, 1994
+	EQ 3MG BASE/ML **	N020239 001	Dec 29, 1993
+	EQ 4MG BASE/4ML (EQ 1MG BASE/ML) **	N020239 002	Mar 11, 1994

SOLUTION; ORAL

GRANISOL

INTRA SANA LABS	EQ 2MG BASE/10ML	A078334 001	Feb 28, 2008
-----------------	------------------	-------------	--------------

KYTRIL

+ ROCHE	EQ 2MG BASE/10ML **	N021238 001	Jun 27, 2001
---------	---------------------	-------------	--------------

TABLET; ORAL

GRANISETRON HYDROCHLORIDE

AUROBINDO PHARMA USA	EQ 1MG BASE	A078725 001	Jan 30, 2008
BARR	EQ 1MG BASE	A078221 001	Dec 31, 2007
EPIC PHARMA LLC	EQ 1MG BASE	A078260 001	Dec 31, 2007
HIKMA	EQ 1MG BASE	A077842 001	Dec 31, 2007
TEVA PHARMS	EQ 1MG BASE	A078080 001	Dec 31, 2007

KYTRIL

+ ROCHE	EQ 1MG BASE **	N020305 001	Mar 16, 1995
+	EQ 2MG BASE **	N020305 002	Jun 15, 1998

GREPAFLOXACIN HYDROCHLORIDE

TABLET; ORAL

RAXAR

OTSUKA	EQ 200MG BASE	N020695 001	Nov 06, 1997
	EQ 400MG BASE	N020695 002	May 14, 1998
	EQ 600MG BASE	N020695 003	May 14, 1998

GRISEOFULVIN, MICROCRYSTALLINE

CAPSULE; ORAL

GRISACTIN

WYETH AYERST	125MG	N050051 002	
	250MG	N050051 001	

SUSPENSION; ORAL

GRIFULVIN V

+ JOHNSON AND JOHNSON	125MG/5ML **	N050448 001	
-----------------------	--------------	-------------	--

TABLET; ORAL

GRIFULVIN V

J AND J	125MG	A060618 001	
	250MG	A060618 002	
	500MG	A060618 003	
VALEANT LUXEMBOURG	125MG	A062279 001	
	250MG **	A062279 002	

GRISACTIN

WYETH AYERST	500MG	A060212 001	
--------------	-------	-------------	--

GRISEOFULVIN, MICROSIZE

SUSPENSION; ORAL

GRIFULVIN V

VALEANT LUXEMBOURG	125MG/5ML **	A062483 001	Jan 26, 1984
--------------------	--------------	-------------	--------------

TABLET; ORAL

GRIFULVIN V

VALEANT LUXEMBOURG	500MG	A062279 003	
--------------------	-------	-------------	--

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

GRISEOFULVIN, ULTRAMICROCRYSTALLINE

TABLET; ORAL

GRISACTIN ULTRA

WYETH AYERST

125MG

A062178 001

165MG

A062438 001 Nov 17, 1983

250MG

A062178 002

330MG

A062438 002 Nov 17, 1983

ULTRAGRIS-165

PLIVA

165MG

A062645 001 Jun 30, 1992

ULTRAGRIS-330

PLIVA

330MG

A062646 001 Jun 30, 1992

GUAIFENESIN; HYDROCODONE BITARTRATE

SOLUTION; ORAL

FLOWTUSS

CHARTWELL RX

200MG/5ML; 2.5MG/5ML

N022424 001 May 14, 2015

OBREDON

+ SOVEREIGN PHARMS

200MG/5ML; 2.5MG/5ML

N205474 001 Nov 14, 2014

TABLET; ORAL

XTRELUS

+ ECI PHARMS LLC

400MG; 5MG

N208085 001 Apr 25, 2018

GUAIFENESIN; HYDROCODONE BITARTRATE; PSEUDOEPHEDRINE HYDROCHLORIDE

SOLUTION; ORAL

HYCOFENIX

+ CHARTWELL RX

200MG/5ML; 2.5MG/5ML; 30MG/5ML

N022279 001 May 14, 2015

GUAIFENESIN; PSEUDOEPHEDRINE HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL

GUAIFENESIN AND PSEUDOEPHEDRINE HYDROCHLORIDE

GRANULES

600MG; 60MG

A216082 001 Aug 22, 2022

1.2GM; 120MG

A216082 002 Aug 22, 2022

GUANABENZ ACETATE

TABLET; ORAL

GUANABENZ ACETATE

ANI PHARMS

EQ 4MG BASE

A074149 001 Apr 07, 1995

EQ 4MG BASE

A074267 001 Jun 01, 1994

EQ 8MG BASE

A074149 002 Apr 07, 1995

EQ 8MG BASE

A074267 002 Jun 01, 1994

CHARTWELL RX

EQ 4MG BASE

A074517 001 Sep 30, 1998

EQ 8MG BASE

A074517 002 Sep 30, 1998

WATSON LABS

EQ 4MG BASE

A074025 001 Feb 28, 1994

EQ 8MG BASE

A074025 002 Feb 28, 1994

WYTENSIN

WYETH AYERST

EQ 4MG BASE

N018587 001 Sep 07, 1982

EQ 8MG BASE

N018587 002 Sep 07, 1982

EQ 16MG BASE

N018587 003 Sep 07, 1982

GUANADREL SULFATE

TABLET; ORAL

HYLOREL

PHARMACIA AND UPJOHN

10MG

N018104 001 Dec 29, 1982

25MG

N018104 002 Dec 29, 1982

GUANETHIDINE MONOSULFATE

TABLET; ORAL

GUANETHIDINE MONOSULFATE

WATSON LABS

EQ 10MG SULFATE

A086113 001 Mar 26, 1985

EQ 25MG SULFATE

A086114 001 Mar 26, 1985

ISMELIN

NOVARTIS

EQ 10MG SULFATE

N012329 001

EQ 25MG SULFATE

N012329 002

GUANETHIDINE MONOSULFATE; HYDROCHLOROTHIAZIDE

TABLET; ORAL

ESIMIL

NOVARTIS

10MG; 25MG

N013553 001

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

GUANFACINE HYDROCHLORIDE

TABLET;ORAL

GUANFACINE HYDROCHLORIDE

WATSON LABS

EQ 1MG BASE

A074762 001 Jun 25, 1997

EQ 2MG BASE

A074762 002 Jun 25, 1997

TENEX

+ PROMIUS PHARMA

EQ 1MG BASE **

N019032 001 Oct 27, 1986

+

EQ 2MG BASE **

N019032 002 Nov 07, 1988

EQ 3MG BASE **

N019032 003 Nov 07, 1988

TABLET, EXTENDED RELEASE;ORAL

GUANFACINE HYDROCHLORIDE

IMPAX LABS INC

EQ 1MG BASE

A202238 001 Oct 20, 2015

EQ 2MG BASE

A202238 002 Oct 20, 2015

EQ 3MG BASE

A202238 003 Oct 20, 2015

EQ 4MG BASE

A202238 004 Oct 20, 2015

NORVIUM BIOSCIENCE

EQ 1MG BASE

A202578 001 Jun 02, 2015

EQ 2MG BASE

A202578 002 Jun 02, 2015

EQ 3MG BASE

A202578 003 Jun 02, 2015

EQ 4MG BASE

A202578 004 Jun 02, 2015

WANBANG BIOPHARMS

EQ 1MG BASE

A217638 001 Jun 12, 2024

EQ 2MG BASE

A217638 002 Jun 12, 2024

EQ 3MG BASE

A217638 003 Jun 12, 2024

EQ 4MG BASE

A217638 004 Jun 12, 2024

GUANIDINE HYDROCHLORIDE

TABLET;ORAL

GUANIDINE HYDROCHLORIDE

MERCK SHARP DOHME

125MG

N001546 001

HALAZEPAM

TABLET;ORAL

PAXIPAM

SCHERING

20MG

N017736 003

40MG

N017736 004

HALCINONIDE

CREAM;TOPICAL

HALOG

WESTWOOD SQUIBB

0.025%

N017818 001

HALOG-E

SUN PHARM INDS INC

0.1%

N018234 001

OINTMENT;TOPICAL

HALOG

BRISTOL MYERS SQUIBB

0.025%

N018125 001

+ SUN PHARM INDS INC

0.1%

N017824 001

SOLUTION;TOPICAL

HALOG

+ SUN PHARM INDS INC

0.1% **

N017823 001

HALOBETASOL PROPIONATE

CREAM;TOPICAL

ULTRAVATE

+ SUN PHARM INDS INC

0.05% **

N019967 001 Dec 27, 1990

LOTION;TOPICAL

HALOBETASOL PROPIONATE

PADAGIS ISRAEL

0.05%

A211464 001 Jun 03, 2020

ULTRAVATE

+ MICAL PHARMS

0.05%

N208183 001 Nov 06, 2015

OINTMENT;TOPICAL

HALOBETASOL PROPIONATE

COSETTE

0.05%

A077721 001 Sep 07, 2006

ENCUBE

0.05%

A209978 001 Mar 20, 2018

FOUGERA PHARMS

0.05%

A076903 001 Dec 16, 2004

ULTRAVATE

+ SUN PHARM INDS INC

0.05% **

N019968 001 Dec 17, 1990

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

HALOFANTRINE HYDROCHLORIDE

TABLET;ORAL

HALFAN

GLAXOSMITHKLINE 250MG N020250 001 Jul 24, 1992

HALOPERIDOL

TABLET;ORAL

HALDOL

+	ORTHO MCNEIL	0.5MG **	N015921 001
+		1MG **	N015921 002
+		2MG **	N015921 003
+		5MG **	N015921 004
+		10MG **	N015921 005
+		20MG **	N015921 006 Feb 02, 1982

HALDOL SOLUTAB

ORTHO MCNEIL PHARM 1MG N017079 001

HALOPERIDOL

	ACTAVIS GROUP	0.5MG	A200854 001 Jul 01, 2022
		1MG	A200854 002 Jul 01, 2022
		2MG	A200854 003 Jul 01, 2022
		5MG	A200854 004 Jul 01, 2022
		10MG	A200854 005 Jul 01, 2022
		20MG	A200854 006 Jul 01, 2022
	AIPING PHARM INC	0.5MG	A071128 001 Feb 17, 1987
		1MG	A071129 001 Feb 17, 1987
		20MG	A071133 001 May 12, 1987
	CHARTWELL RX	2MG	A071209 004 Nov 17, 1986
	DURAMED PHARMS BARR	0.5MG	A071216 001 Dec 04, 1986
		1MG	A071217 001 Dec 04, 1986
		2MG	A071218 001 Dec 04, 1986
		5MG	A071219 001 Dec 04, 1986
		10MG	A071220 001 Jul 07, 1987
		20MG	A071221 001 Jul 07, 1987
	LEDERLE	0.5MG	A072727 001 Sep 19, 1989
		1MG	A072728 001 Sep 19, 1989
		2MG	A072729 001 Sep 19, 1989
		5MG	A072730 001 Sep 19, 1989
		10MG	A072731 001 Sep 19, 1989
		20MG	A072732 001 Sep 19, 1989
	PAR PHARM	20MG	A071328 001 Jul 20, 1987
	PUREPAC PHARM	0.5MG	A071071 001 Nov 03, 1986
		1MG	A071072 001 Nov 03, 1986
		2MG	A071073 001 Nov 03, 1986
		5MG	A071074 001 Nov 03, 1986
		10MG	A071075 001 Aug 04, 1987
		20MG	A071076 001 Aug 04, 1987
	QUANTUM PHARMICS	0.5MG	A071255 001 Feb 17, 1987
		1MG	A071269 001 Feb 17, 1987
		2MG	A071256 001 Feb 17, 1987
		5MG	A071257 001 Feb 17, 1987
	ROYCE LABS	0.5MG	A071722 001 Dec 24, 1987
		1MG	A071723 001 Dec 24, 1987
		2MG	A071724 001 Dec 24, 1987
		5MG	A071725 001 Dec 24, 1987
		10MG	A072121 001 Dec 24, 1987
		20MG	A072122 001 Dec 24, 1987
	SCS	0.5MG	A070720 001 Jun 10, 1986
		1MG	A070721 001 Jun 10, 1986
		2MG	A070722 001 Jun 10, 1986
		5MG	A070723 001 Jun 10, 1986
		10MG	A070724 001 Jun 10, 1986
		20MG	A070725 001 Sep 24, 1986
	STRIDES PHARMA	0.5MG	A071235 002 Nov 03, 1986
		1MG	A071235 003 Nov 03, 1986
		2MG	A071235 001 Nov 03, 1986
		5MG	A071235 004 Nov 03, 1986
		10MG	A071235 005 Jul 20, 1987
	WATSON LABS	0.5MG	A070981 001 Mar 06, 1987
		0.5MG	A071571 001 Jun 03, 1988
		1MG	A070982 001 Mar 06, 1987

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

HALOPERIDOL

TABLET; ORAL

HALOPERIDOL

1MG	A071572 001	Jun 03, 1988
2MG	A070983 001	Mar 06, 1987
2MG	A071573 001	Jun 03, 1988
5MG	A070984 001	Mar 06, 1987
5MG	A071374 001	Jun 03, 1988
10MG	A071375 001	Jun 03, 1988
10MG	A072113 001	Aug 27, 1991
20MG	A071376 001	Jun 03, 1988
20MG	A072353 001	Aug 27, 1991

HALOPERIDOL DECANOATE

INJECTABLE; INJECTION

HALOPERIDOL DECANOATE

HOSPIRA

EQ 50MG BASE/ML

A075176 001 Feb 09, 2000

EQ 100MG BASE/ML

A075176 002 Feb 09, 2000

SANDOZ

EQ 50MG BASE/ML

A076463 001 Jun 24, 2005

EQ 100MG BASE/ML

A076463 002 Jun 24, 2005

TEVA PHARMS USA

EQ 50MG BASE/ML

A075393 001 May 11, 1999

EQ 100MG BASE/ML

A075393 002 May 11, 1999

HALOPERIDOL LACTATE

CONCENTRATE; ORAL

HALDOL

+ ORTHO MCNEIL

EQ 2MG BASE/ML **

N015922 001

HALOPERIDOL

ALPHARMA

EQ 2MG BASE/ML

A070318 001 Apr 11, 1986

MORTON GROVE

EQ 2MG BASE/ML

A070710 001 Mar 07, 1986

SCS

EQ 2MG BASE/ML

A070726 001 Jun 10, 1986

TEVA

EQ 2MG BASE/ML

A071015 001 Aug 25, 1987

TEVA PHARMS

EQ 2MG BASE/ML

A071617 001 Dec 01, 1988

HALOPERIDOL INTENSOL

HIKMA

EQ 2MG BASE/ML

A072045 001 Apr 12, 1988

INJECTABLE; INJECTION

HALDOL

+ JANSSEN PHARMS

EQ 5MG BASE/ML **

N015923 001

HALOPERIDOL

ABRAXIS PHARM

EQ 5MG BASE/ML

A071187 001 Jan 20, 1987

BAXTER HLTHCARE CORP

EQ 5MG BASE/ML

A076791 001 Aug 25, 2004

EQ 5MG BASE/ML

A076828 001 Aug 25, 2004

EPIC PHARMA LLC

EQ 5MG BASE/ML

A204849 001 Sep 06, 2017

FOSUN PHARMA

EQ 5MG BASE/ML

A076464 001 Sep 29, 2004

MARSAM PHARMS LLC

EQ 5MG BASE/ML

A072516 001 Feb 25, 1993

EQ 5MG BASE/ML

A072517 001 Feb 25, 1993

SMITH AND NEPHEW

EQ 5MG BASE/ML

A070802 001 Dec 14, 1987

SOLOPAK

EQ 5MG BASE/ML

A070800 001 Dec 14, 1987

EQ 5MG BASE/ML

A070801 001 Dec 14, 1987

EQ 5MG BASE/ML

A070864 001 Dec 14, 1987

TEVA PHARMS USA

EQ 5MG BASE/ML

A076035 001 Aug 29, 2001

WATSON LABS

EQ 5MG BASE/ML

A070713 001 May 17, 1988

EQ 5MG BASE/ML

A070714 001 May 17, 1988

EQ 5MG BASE/ML

A070744 001 May 17, 1988

SOLUTION; ORAL

HALOPERIDOL LACTATE

ACTAVIS MID ATLANTIC

EQ 1MG BASE/ML

A074536 001 Nov 02, 1995

HALOPROGIN

CREAM; TOPICAL

HALOTEX

WESTWOOD SQUIBB

1%

N016942 001

SOLUTION; TOPICAL

HALOTEX

WESTWOOD SQUIBB

1%

N016943 001

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

HALOTHANE

LIQUID; INHALATION

FLUOTHANE

WYETH AYERST 99.99% N011338 001

HALOTHANE

BH 99.99% A084977 001

HALOCARBON 99.99% A080810 001

HOSPIRA 99.99% A083254 001

HEPARIN CALCIUM

INJECTABLE; INJECTION

CALCIPARINE

SANOFI AVENTIS US 25,000 UNITS/ML N018237 001

HEPARIN SODIUM

INJECTABLE; INJECTION

HEPARIN LOCK FLUSH

HOSPIRA 100 UNITS/ML N005264 010

INTL MEDICATION 10 UNITS/ML A086357 001

500 UNITS/ML A086357 002

LUITPOLD 10 UNITS/ML A089063 001

Oct 09, 1985

100 UNITS/ML A089064 001

Oct 09, 1985

PARKE DAVIS 10 UNITS/ML N017346 006

SMITH AND NEPHEW 10 UNITS/ML A087904 001

Apr 20, 1983

10 UNITS/ML A087958 001

Apr 20, 1983

10 UNITS/ML A088458 001

Jul 26, 1984

10 UNITS/ML A088580 001

Oct 25, 1984

100 UNITS/ML A087906 001

Apr 20, 1983

100 UNITS/ML A087959 001

Apr 20, 1983

100 UNITS/ML A088460 001

Jul 26, 1984

100 UNITS/ML A088581 001

Oct 25, 1984

SOLOPAK 10 UNITS/ML A087903 001

Apr 20, 1983

10 UNITS/ML A088457 001

Oct 25, 1984

100 UNITS/ML A087905 001

Apr 20, 1983

100 UNITS/ML A088459 001

Jul 26, 1984

HEPARIN SODIUM

ABRAXIS PHARM 1,000 UNITS/ML N017033 001

1,000 UNITS/ML N017979 001

5,000 UNITS/ML N017979 003

10,000 UNITS/ML N017979 002

ASPEN GLOBAL INC 1,000 UNITS/ML N000552 008

5,000 UNITS/ML N000552 009

10,000 UNITS/ML N000552 010

CASI PHARMS INC 5,000 UNITS/ML A091659 001 Jun 08, 2011

CHAMBERLIN PARENTERL 1,000 UNITS/ML N017130 001

5,000 UNITS/ML N017130 002

10,000 UNITS/ML N017130 003

20,000 UNITS/ML N017130 004

DELL LABS 1,000 UNITS/ML N017540 001

5,000 UNITS/ML N017540 002

10,000 UNITS/ML N017540 003

20,000 UNITS/ML N017540 004

40,000 UNITS/ML N017540 005

DR REDDYS 1,000 UNITS/ML A040007 001 Jun 07, 1996

1,000 UNITS/ML N017064 002

2,500 UNITS/ML N017064 015

3,000 UNITS/ML N017064 016

4,000 UNITS/ML N017064 017

5,000 UNITS/ML N017064 003

6,000 UNITS/ML N017064 018

7,500 UNITS/ML N017064 019

10,000 UNITS/ML N017064 004

20,000 UNITS/ML N017064 005

40,000 UNITS/ML N017064 006

EPIC PHARMA LLC 1,000 UNITS/ML N017486 001

5,000 UNITS/ML N017486 002

10,000 UNITS/ML N017486 003

20,000 UNITS/ML N017486 004

40,000 UNITS/ML N017486 005

FRESENIUS KABI USA 1,000 UNITS/ML N017651 005

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

HEPARIN SODIUM

INJECTABLE; INJECTION

HEPARIN SODIUM

		5,000 UNITS/ML **	N017029	002	
		10,000 UNITS/ML	N017651	003	
		20,000 UNITS/ML	N017651	008	
+	HIKMA	1,000 UNITS/ML **	N017007	001	
+		2,500 UNITS/ML **	N017007	007	
+		5,000 UNITS/ML **	N017007	002	
+		5,000 UNITS/0.5ML **	N017007	010	
+		7,500 UNITS/ML **	N017007	003	
+		10,000 UNITS/ML **	N017007	004	
+		15,000 UNITS/ML **	N017007	005	
+		20,000 UNITS/ML **	N017007	006	
	HOSPIRA	2,500 UNITS/ML	A088099	001	Apr 28, 1983
		10,000 UNITS/ML	A040095	001	Jul 26, 1996
	LILLY	1,000 UNITS/ML	N005521	001	
		10,000 UNITS/ML	N005521	002	
		20,000 UNITS/ML	N005521	004	
	LUITPOLD	1,000 UNITS/ML	A087452	001	Oct 31, 1983
	PARKE DAVIS	1,000 UNITS/ML	N017346	001	
		5,000 UNITS/ML	N017346	002	
		7,500 UNITS/ML	N017346	003	
		10,000 UNITS/ML	N017346	004	
		20,000 UNITS/ML	N017346	005	
+	PFIZER	10,000 UNITS/ML	N201370	003	Jul 21, 2011
	PHARM SPEC	1,000 UNITS/ML	N017780	001	
		5,000 UNITS/ML	N017780	002	
		10,000 UNITS/ML	N017780	003	
		20,000 UNITS/ML	N017780	004	
		40,000 UNITS/ML	N017780	005	
	PHARMACIA AND UPJOHN	1,000 UNITS/ML	N004570	001	
		5,000 UNITS/ML	N004570	002	
		10,000 UNITS/ML	N004570	003	
	SMITH AND NEPHEW	1,000 UNITS/ML	A088239	001	Jul 26, 1984
	SOLOPAK	1,000 UNITS/ML	A087043	001	
		5,000 UNITS/ML	A087077	001	
		5,000 UNITS/0.5ML	A087395	001	
		10,000 UNITS/ML	A087107	001	
		10,000 UNITS/0.5ML	A087363	001	
	WATSON LABS INC	1,000 UNITS/ML	A040008	001	Oct 10, 1995
	HEPARIN SODIUM 1,000 UNITS	IN DEXTROSE 5% IN PLASTIC CONTAINER			
	MCGAW	200 UNITS/100ML	N019130	001	Dec 31, 1984
	HEPARIN SODIUM 1,000 UNITS	IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER			
	B BRAUN	200 UNITS/100ML	N019042	001	Mar 29, 1985
	HEPARIN SODIUM 10,000 UNITS	AND DEXTROSE 5% IN PLASTIC CONTAINER			
	BAXTER HLTHCARE	2,000 UNITS/100ML	N018814	002	Jul 09, 1985
	HEPARIN SODIUM 10,000 UNITS	IN DEXTROSE 5%			
	HOSPIRA	10,000 UNITS/100ML	N018911	006	Jan 30, 1985
	HEPARIN SODIUM 10,000 UNITS	IN SODIUM CHLORIDE 0.45%			
	HOSPIRA	10,000 UNITS/100ML	N018911	001	Jan 30, 1985
		10,000 UNITS/100ML	N018916	005	Jan 31, 1984
	HEPARIN SODIUM 10,000 UNITS	IN SODIUM CHLORIDE 0.9%			
	HOSPIRA	10,000 UNITS/100ML	N018911	003	Jan 30, 1985
		10,000 UNITS/100ML	N018916	002	Jan 31, 1984
	HEPARIN SODIUM 12,500 UNITS	IN DEXTROSE 5%			
	HOSPIRA	5,000 UNITS/100ML	N018911	007	Jan 30, 1985
	HEPARIN SODIUM 12,500 UNITS	IN SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER			
	B BRAUN	5,000 UNITS/100ML	N019802	001	Jul 20, 1992
	HEPARIN SODIUM 12,500 UNITS	IN SODIUM CHLORIDE 0.9%			
	HOSPIRA	5,000 UNITS/100ML	N018911	005	Jan 30, 1985
		5,000 UNITS/100ML	N018916	003	Jan 31, 1984
	HEPARIN SODIUM 2,000 UNITS	IN DEXTROSE 5% IN PLASTIC CONTAINER			
	MCGAW	200 UNITS/100ML	N019130	003	Dec 31, 1984
	HEPARIN SODIUM 2,000 UNITS	IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER			
	B BRAUN	200 UNITS/100ML	N019042	002	Mar 29, 1985
	HEPARIN SODIUM 20,000 UNITS	AND DEXTROSE 5% IN PLASTIC CONTAINER			
	BAXTER HLTHCARE	4,000 UNITS/100ML	N018814	001	Oct 31, 1983

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

HEPARIN SODIUM

INJECTABLE; INJECTION

HEPARIN SODIUM 20,000 UNITS	IN DEXTROSE 5%				
+ FRESENIUS KABI USA	40 UNITS/ML	N017029	021	Aug 24,	2017
HEPARIN SODIUM 20,000 UNITS	IN DEXTROSE 5% IN PLASTIC CONTAINER				
HOSPIRA	4,000 UNITS/100ML	N019805	001	Jan 25,	1989
HEPARIN SODIUM 25,000 UNITS	AND DEXTROSE 5% IN PLASTIC CONTAINER				
BAXTER HLTHCARE	5,000 UNITS/100ML	N018814	003	Jul 09,	1985
	10,000 UNITS/100ML	N018814	004	Jul 02,	1987
HEPARIN SODIUM 25,000 UNITS	IN DEXTROSE 5%				
HOSPIRA	5,000 UNITS/100ML	N018911	009	Jan 30,	1985
	10,000 UNITS/100ML	N018911	008	Jan 30,	1985
HEPARIN SODIUM 25,000 UNITS	IN DEXTROSE 5% IN PLASTIC CONTAINER				
B BRAUN	5,000 UNITS/100ML	N019134	001	Mar 29,	1985
HOSPIRA	5,000 UNITS/100ML	N019805	002	Jan 25,	1989
HEPARIN SODIUM 25,000 UNITS	IN SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER				
B BRAUN	5,000 UNITS/100ML	N019802	005	Jul 20,	1992
	10,000 UNITS/100ML	N019802	002	Jul 20,	1992
HEPARIN SODIUM 25,000 UNITS	IN SODIUM CHLORIDE 0.9%				
HOSPIRA	5,000 UNITS/100ML	N018911	004	Jan 30,	1985
HEPARIN SODIUM 25,000 UNITS	IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER				
B BRAUN	5,000 UNITS/100ML	N019135	001	Mar 29,	1985
	5,000 UNITS/100ML	N019802	003	Jul 20,	1992
HOSPIRA	5,000 UNITS/100ML	N018916	009	Jan 31,	1984
HEPARIN SODIUM 5,000 UNITS	AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER				
BAXTER HLTHCARE	500 UNITS/100ML	N018609	003	Apr 28,	1982
HEPARIN SODIUM 5,000 UNITS	IN DEXTROSE 5% IN PLASTIC CONTAINER				
MCGAW	1,000 UNITS/100ML	N019130	002	Dec 31,	1984
HEPARIN SODIUM 5,000 UNITS	IN SODIUM CHLORIDE 0.45%				
HOSPIRA	100 UNITS/ML	N018911	002	Jan 30,	1985
	100 UNITS/ML	N018916	004	Jan 31,	1984
HEPARIN SODIUM 5,000 UNITS	IN SODIUM CHLORIDE 0.9%				
HOSPIRA	1,000 UNITS/100ML	N018916	001	Jan 31,	1984
HEPARIN SODIUM 5,000 UNITS	IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER				
B BRAUN	1,000 UNITS/100ML	N019042	004	Mar 29,	1985
HEPARIN SODIUM PRESERVATIVE FREE					
DR REDDYS	1,000 UNITS/ML	A089464	001	Jun 03,	1986
HOSPIRA	2,000 UNITS/ML	N005264	013	Apr 07,	1986
	2,500 UNITS/ML	N005264	014	Apr 07,	1986
PHARMA SERVE NY	1,000 UNITS/ML	A086129	001		
LIPO-HEPIN					
3M	1,000 UNITS/0.5ML	N017027	001		
	1,000 UNITS/ML	N017027	006		
	5,000 UNITS/0.5ML	N017027	002		
	5,000 UNITS/ML	N017027	008		
	7,500 UNITS/0.5ML	N017027	010		
	10,000 UNITS/0.5ML	N017027	003		
	10,000 UNITS/ML	N017027	009		
	15,000 UNITS/0.5ML	N017027	011		
	20,000 UNITS/0.5ML	N017027	004		
	20,000 UNITS/ML	N017027	007		
	40,000 UNITS/ML	N017027	005		
LIQUAEMIN LOCK FLUSH					
ASPEN GLOBAL INC	100 UNITS/ML	N000552	007		
LIQUAEMIN SODIUM					
ASPEN GLOBAL INC	1,000 UNITS/ML	N000552	004		
	5,000 UNITS/ML	N000552	003		
	10,000 UNITS/ML	N000552	005		
	20,000 UNITS/ML	N000552	001		
	40,000 UNITS/ML	N000552	002		
LIQUAEMIN SODIUM PRESERVATIVE FREE					
ASPEN GLOBAL INC	1,000 UNITS/ML	N000552	011	Apr 11,	1986
	5,000 UNITS/ML	N000552	012	Apr 11,	1986
	10,000 UNITS/ML	N000552	013	Apr 11,	1986
PANHEPRIN					
HOSPIRA	1,000 UNITS/ML	N005264	004		
	5,000 UNITS/ML	N005264	006		
	10,000 UNITS/ML	N005264	007		
	20,000 UNITS/ML	N005264	008		

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

HEPARIN SODIUM

INJECTABLE; INJECTION

PANHEPRIN

40,000 UNITS/ML

N005264 009

SODIUM HEPARIN

ABRAXIS PHARM

5,000 UNITS/ML

N017033 002

10,000 UNITS/ML

N017033 003

20,000 UNITS/ML

N017033 004

BAXTER HLTHCARE

1,000 UNITS/ML

N017036 001 Mar 04, 1988

HETACILLIN

FOR SUSPENSION; ORAL

VERSAPEN

BRISTOL

EQ 112.5MG AMPICIL/ML

A061398 001

EQ 112.5MG AMPICIL/5ML

N050060 001

EQ 112.5MG AMPICIL/ML

N050060 003

EQ 225MG AMPICIL/5ML

A061398 002

HETACILLIN POTASSIUM

CAPSULE; ORAL

VERSAPEN-K

BRISTOL

EQ 225MG AMPICIL

A061396 001

EQ 450MG AMPICIL

A061396 002

HEXACHLOROPHENE

AEROSOL; TOPICAL

SEPTISOL

VESTAL LABS

0.23%

N017424 001

TURGEX

XTTRIUM

3%

N018375 001

EMULSION; TOPICAL

HEXA-GERM

HUNTINGTON LABS

3%

N017411 001

PHISOHEX

SANOFI AVENTIS US

3%

N006882 001

3%

N008402 001

SOY-DOME

BAYER PHARMS

3%

N017405 001

TURGEX

XTTRIUM

3%

N019055 001 Nov 30, 1984

SOAP; TOPICAL

GAMOPHEN

ARBROOK

2%

N006270 003

SOLUTION; TOPICAL

DIAL

DIAL

0.25%

N017421 002

GERMA-MEDICA

HUNTINGTON LABS

1%

N017412 001

GERMA-MEDICA "MG"

HUNTINGTON LABS

0.25%

N017412 002

SEPTI-SOFT

CALGON

0.25%

N017460 001

SEPTISOL

VESTAL LABS

0.25%

N017423 001

SPONGE; TOPICAL

E-Z SCRUB

BECTON DICKINSON

450MG

N017452 001

HEXASCRUB

PROF DSPLS

3%

N018363 001

PHISO-SCRUB

SANOFI AVENTIS US

3%

N017446 001

SCRUBTEAM SURGICAL SPONGEBRUSH

3M

330MG

N017413 001

HEXAFLUORENIUM BROMIDE

INJECTABLE; INJECTION

MYLAXEN

MEDPOINTE PHARM HLC

20MG/ML

N009789 003

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

HEXOCYCLIUM METHYLSULFATE

TABLET; ORAL

TRAL

ABBVIE	25MG	N010599	001
--------	------	---------	-----

HEXYLCAINE HYDROCHLORIDE

SOLUTION; TOPICAL

CYCLAINE

MERCK	5%	N008472	001
-------	----	---------	-----

HISTAMINE PHOSPHATE

INJECTABLE; INJECTION

HISTAMINE PHOSPHATE

LILLY	EQ 0.1MG BASE/ML	N000734	003
-------	------------------	---------	-----

	EQ 0.2MG BASE/ML	N000734	002
--	------------------	---------	-----

	EQ 1MG BASE/ML	N000734	001
--	----------------	---------	-----

HISTRELIN ACETATE

IMPLANT; SUBCUTANEOUS

VANTAS

+ ENDO OPERATIONS	50MG	N021732	001	Oct 12, 2004
-------------------	------	---------	-----	--------------

INJECTABLE; INJECTION

SUPPRELIN

SHIRE	EQ 0.2MG BASE/ML	N019836	001	Dec 24, 1991
-------	------------------	---------	-----	--------------

	EQ 0.5MG BASE/ML	N019836	002	Dec 24, 1991
--	------------------	---------	-----	--------------

	EQ 1MG BASE/ML	N019836	003	Dec 24, 1991
--	----------------	---------	-----	--------------

HOMATROPINE METHYLBROMIDE

TABLET; ORAL

HOMAPIN-10

MISSION PHARMA	10MG	A086308	001
----------------	------	---------	-----

HOMAPIN-5

MISSION PHARMA	5MG	A086309	001
----------------	-----	---------	-----

TABLET, CHEWABLE; ORAL

EQUIPIN

MISSION PHARMA	3MG	A086310	001
----------------	-----	---------	-----

HOMATROPINE METHYLBROMIDE; HYDROCODONE BITARTRATE

SYRUP; ORAL

HYDROCODONE BITARTRATE AND HOMATROPINE METHYLBROMIDE

GENUS	1.5MG/5ML; 5MG/5ML	A040613	001	Feb 08, 2008
-------	--------------------	---------	-----	--------------

IVAX SUB TEVA PHARMS	1.5MG/5ML; 5MG/5ML	A040285	001	Jul 19, 1999
----------------------	--------------------	---------	-----	--------------

NOSTRUM LABS INC	1.5MG/5ML; 5MG/5ML	A210663	001	Jun 11, 2019
------------------	--------------------	---------	-----	--------------

SCIEGEN PHARMS INC	1.5MG/5ML; 5MG/5ML	A204765	001	Mar 06, 2017
--------------------	--------------------	---------	-----	--------------

HYDROPANE

HALSEY	1.5MG/5ML; 5MG/5ML	A088066	001	Jun 28, 1985
--------	--------------------	---------	-----	--------------

TABLET; ORAL

HOMATROPINE METHYLBROMIDE AND HYDROCODONE BITARTRATE

ACTAVIS ELIZABETH	1.5MG; 5MG	A040295	001	Dec 01, 2000
-------------------	------------	---------	-----	--------------

NOVEL LABS INC	1.5MG; 5MG	A091528	001	Apr 20, 2011
----------------	------------	---------	-----	--------------

TUSSIGON

KING PHARMS	1.5MG; 5MG	A088508	001	Jul 30, 1985
-------------	------------	---------	-----	--------------

HYDRALAZINE HYDROCHLORIDE

INJECTABLE; INJECTION

APRESOLINE

+ NOVARTIS	20MG/ML **	N008303	003
------------	------------	---------	-----

HYDRALAZINE HYDROCHLORIDE

ABRAXIS PHARM	20MG/ML	A089532	001	Aug 11, 1987
---------------	---------	---------	-----	--------------

NORVIUM BIOSCIENCE	20MG/ML	A204680	001	Apr 28, 2016
--------------------	---------	---------	-----	--------------

SMITH AND NEPHEW	20MG/ML	A088518	001	Apr 20, 1984
------------------	---------	---------	-----	--------------

SOLOPAK	20MG/ML	A088517	001	Aug 22, 1985
---------	---------	---------	-----	--------------

TEVA PARENTERAL	20MG/ML	A040373	001	Feb 23, 2000
-----------------	---------	---------	-----	--------------

TABLET; ORAL

APRESOLINE

+ NOVARTIS	10MG **	N008303	004
------------	---------	---------	-----

	25MG **	N008303	001
--	---------	---------	-----

	50MG **	N008303	002
--	---------	---------	-----

	100MG **	N008303	005
--	----------	---------	-----

DRALZINE

TEVA	25MG	A084301	001
------	------	---------	-----

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

HYDRALAZINE HYDROCHLORIDE

TABLET;ORAL

HYDRALAZINE HYDROCHLORIDE

ACTAVIS ELIZABETH	25MG	A088560	001	Oct 04, 1984
	50MG	A088649	001	Oct 18, 1984
ACTAVIS GRP PTC	10MG	A091679	001	Mar 04, 2013
	25MG	A091679	002	Mar 04, 2013
	50MG	A091679	003	Mar 04, 2013
	100MG	A091679	004	Mar 04, 2013
ANDA REPOSITORY	10MG	A089359	001	Jul 25, 1986
	25MG	A089258	001	May 05, 1986
	50MG	A089259	001	May 05, 1986
	100MG	A088729	001	Apr 11, 1985
ASCOT	25MG	A088310	001	Dec 19, 1984
	50MG	A088311	001	Dec 19, 1984
CHARTWELL RX	10MG	A088846	001	Feb 26, 1985
	25MG	A088847	001	Feb 26, 1985
	50MG	A088848	001	Feb 26, 1985
	100MG	A088849	001	Feb 26, 1985
HALSEY	10MG	A089218	001	Jan 22, 1986
	25MG	A089130	001	Jan 15, 1986
	50MG	A089222	001	Jan 22, 1986
	100MG	A089178	001	Jan 15, 1986
IMPAX LABS	25MG	A084922	001	
	50MG	A084923	001	
INVAGEN PHARMS	10MG	A090255	001	Dec 15, 2008
	25MG	A090255	002	Dec 15, 2008
	50MG	A090255	003	Dec 15, 2008
	100MG	A090255	004	Dec 15, 2008
IVAX SUB TEVA PHARMS	10MG	A084443	001	
	25MG	A084437	001	
	50MG	A084469	002	
	100MG	A084581	001	
MUTUAL PHARM	10MG	A088728	001	Apr 11, 1985
	25MG	A084106	002	
	50MG	A084107	002	
NORVIUM BIOSCIENCE	10MG	A090413	001	Dec 08, 2010
	25MG	A090413	002	Dec 08, 2010
	50MG	A090413	003	Dec 08, 2010
	100MG	A090413	004	Dec 08, 2010
PUREPAC PHARM	25MG	A088177	001	Jul 29, 1983
	50MG	A088178	001	Aug 15, 1983
QUANTUM PHARMICS	10MG	A088671	001	May 01, 1984
	25MG	A088657	001	Jun 15, 1984
	50MG	A088652	001	May 08, 1984
	100MG	A088686	001	May 01, 1984
RISING	10MG	A209251	001	Jul 09, 2018
	25MG	A209251	002	Jul 09, 2018
	50MG	A209251	003	Jul 09, 2018
	100MG	A209251	004	Jul 09, 2018
STRIDES PHARMA	10MG	A200770	004	Jun 25, 2019
SUPERPHARM	10MG	A088787	001	Aug 28, 1984
	25MG	A088788	001	Aug 28, 1984
	50MG	A088789	001	Aug 28, 1984
UPSHER SMITH LABS	10MG	A083241	001	
	25MG	A083560	001	
	50MG	A083561	001	
	50MG	A085088	001	
USL PHARMA	25MG	A087780	001	Mar 29, 1982
	50MG	A087751	001	Mar 29, 1982
VANGARD	25MG	A087712	001	
	50MG	A087908	001	May 07, 1982
VITARINE	25MG	A086088	001	
WATSON LABS	25MG	A084504	001	
	25MG	A085532	002	May 24, 1982
	50MG	A084503	001	
	50MG	A085533	002	May 25, 1982
WEST WARD	25MG	A088240	001	May 27, 1983
	50MG	A088241	001	May 27, 1983

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

HYDRALAZINE HYDROCHLORIDE; HYDROCHLOROTHIAZIDE

CAPSULE; ORAL

APRESAZIDE

+	NOVARTIS	25MG;25MG	A084735	001
+		50MG;50MG	A084810	001
		100MG;50MG	A084811	001

HYDRA-ZIDE

	STRIDES PHARMA	100MG;50MG	A088961	001	Oct 21, 1985
--	----------------	------------	---------	-----	--------------

HYDRALAZINE HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE

	SOLVAY	25MG;25MG	A087608	001	Feb 08, 1982
		50MG;50MG	A087213	001	Feb 08, 1982
		100MG;50MG	A087609	001	Feb 08, 1982
	SUPERPHARM	25MG;25MG	A089200	001	Feb 09, 1987
		50MG;50MG	A089201	001	Feb 09, 1987
	WATSON LABS	25MG;25MG	A085457	001	Mar 04, 1982
		50MG;50MG	A085446	001	Mar 04, 1982
		100MG;50MG	A085440	001	Mar 04, 1982

HYDRALAZINE HYDROCHLORIDE W/ HYDROCHLOROTHIAZIDE 100/50

	IVAX PHARMS	100MG;50MG	A088358	001	Apr 10, 1984
--	-------------	------------	---------	-----	--------------

HYDRALAZINE HYDROCHLORIDE W/ HYDROCHLOROTHIAZIDE 25/25

	IVAX PHARMS	25MG;25MG	A088356	001	Apr 10, 1984
--	-------------	-----------	---------	-----	--------------

HYDRALAZINE HYDROCHLORIDE W/ HYDROCHLOROTHIAZIDE 50/50

	IVAX PHARMS	50MG;50MG	A088357	001	Apr 10, 1984
--	-------------	-----------	---------	-----	--------------

TABLET; ORAL

APRESOLINE-ESIDRIX

	NOVARTIS	25MG;15MG	N012026	002
--	----------	-----------	---------	-----

HYDRALAZINE AND HYDROCHLOROTHIAZIDE

	WATSON LABS	25MG;15MG	A085827	001
--	-------------	-----------	---------	-----

HYDROCHLOROTHIAZIDE W/ HYDRALAZINE

	WATSON LABS	25MG;15MG	A085373	001
--	-------------	-----------	---------	-----

HYDRALAZINE HYDROCHLORIDE; HYDROCHLOROTHIAZIDE; RESERPINE

TABLET; ORAL

CAM-AP-ES

	CHARTWELL RX	25MG;15MG;0.1MG	A084897	001
--	--------------	-----------------	---------	-----

HYDRALAZINE HYDROCHLORIDE, HYDROCHLOROTHIAZIDE AND RESERPINE

	IVAX SUB TEVA PHARMS	25MG;15MG;0.1MG	A084291	001
--	----------------------	-----------------	---------	-----

HYDRALAZINE HYDROCHLORIDE-HYDROCHLOROTHIAZIDE-RESERPINE

	MYLAN	25MG;15MG;0.1MG	A087085	001
--	-------	-----------------	---------	-----

HYDRALAZINE, HYDROCHLOROTHIAZIDE W/ RESERPINE

	WATSON LABS	25MG;15MG;0.1MG	A085771	001
--	-------------	-----------------	---------	-----

HYDRAP-ES

	SANDOZ	25MG;15MG;0.1MG	A084876	001
--	--------	-----------------	---------	-----

HYDROCHLOROTHIAZIDE W/ RESERPINE AND HYDRALAZINE

	WATSON LABS	25MG;15MG;0.1MG	A083770	001
--	-------------	-----------------	---------	-----

HYDROSERPINE PLUS (R-H-H)

	IVAX SUB TEVA PHARMS	25MG;15MG;0.1MG	A083877	001
--	----------------------	-----------------	---------	-----

RESERPINE, HYDRALAZINE HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE

	SOLVAY	25MG;15MG;0.1MG	A088376	001	Oct 28, 1983
--	--------	-----------------	---------	-----	--------------

	SUN PHARM INDUSTRIES	25MG;15MG;0.1MG	A088570	001	Apr 10, 1984
--	----------------------	-----------------	---------	-----	--------------

	WATSON LABS	25MG;15MG;0.1MG	A085549	001
--	-------------	-----------------	---------	-----

		25MG;15MG;0.1MG	A087556	001
--	--	-----------------	---------	-----

RESERPINE, HYDROCHLOROTHIAZIDE, AND HYDRALAZINE HYDROCHLORIDE

	LEDERLE	25MG;15MG;0.1MG	A087709	001	May 13, 1982
--	---------	-----------------	---------	-----	--------------

SER-A-GEN

	SOLVAY	25MG;15MG;0.1MG	A087210	001
--	--------	-----------------	---------	-----

SER-AP-ES

	NOVARTIS	25MG;15MG;0.1MG	N012193	005
--	----------	-----------------	---------	-----

UNIPRES

	SOLVAY	25MG;15MG;0.1MG	A085893	001
--	--------	-----------------	---------	-----

		25MG;15MG;0.1MG	A086298	001
--	--	-----------------	---------	-----

HYDRALAZINE HYDROCHLORIDE; RESERPINE

TABLET; ORAL

DRALSERP

	SANDOZ	25MG;0.1MG	A084617	001
--	--------	------------	---------	-----

SERPASIL-APRESOLINE

	NOVARTIS	25MG;0.1MG	N009296	004
--	----------	------------	---------	-----

		50MG;0.2MG	N009296	002
--	--	------------	---------	-----

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

HYDROCHLOROTHIAZIDE

CAPSULE; ORAL

HYDROCHLOROTHIAZIDE

ALEMBIC PHARMS LTD	12.5MG	A200645	001	Nov 30, 2010
APOTEX	12.5MG	A078389	001	May 16, 2008
CHARTWELL MOLECULAR	12.5MG	A091662	001	Jan 27, 2012
HIKMA INTL PHARMS	12.5MG	A077885	001	Nov 26, 2007
IPCA LABS LTD	12.5MG	A079237	001	Apr 02, 2009
IVAX SUB TEVA PHARMS	12.5MG	A077005	001	Jul 13, 2005
NORVIUM BIOSCIENCE	12.5MG	A075640	001	Jan 28, 2000
SUN PHARM INDS INC	12.5MG	A090651	001	Apr 07, 2014

SOLUTION; ORAL

HYDROCHLOROTHIAZIDE

MORTON GROVE	50MG/5ML	A089661	001	Jun 20, 1988
+ ROXANE	50MG/5ML **	A088587	001	Jul 02, 1984

HYDROCHLOROTHIAZIDE INTENSOL

ROXANE	100MG/ML	A088588	001	Jul 02, 1984
--------	----------	---------	-----	--------------

TABLET; ORAL

ESIDRIX

NOVARTIS	25MG	N011793	005	
	50MG	N011793	008	
	100MG	N011793	009	

HYDRO-D

HALSEY	25MG	A086504	001	
	50MG	A083891	002	

HYDROCHLOROTHIAZIDE

ABC HOLDING	50MG	A085672	001	
ACTAVIS ELIZABETH	25MG	A085054	002	
	50MG	A085208	001	
ALRA	25MG	A086369	001	
	50MG	A083554	001	
APOTEX	25MG	A040774	001	Oct 03, 2007
	50MG	A040774	002	Oct 03, 2007
ASCOT	25MG	A087539	001	Feb 03, 1982
	50MG	A087540	001	Feb 03, 1982
AUROLIFE PHARMA LLC	25MG	A083899	001	
	50MG	A085219	001	
BARR	50MG	A084771	001	
CHARTWELL RX	25MG	A085683	001	
	25MG	A087565	001	Mar 09, 1982
	50MG	A083965	001	
	50MG	A084912	001	
DAVA PHARMS INC	100MG	A087060	001	
ELKINS SINN	50MG	A085152	002	
HEATHER	50MG	A084135	001	
HIKMA INTL PHARMS	25MG	A084878	002	Jul 12, 2006
	50MG	A084878	001	
IMPAX LABS	25MG	A084029	001	
	50MG	A083607	002	
	100MG	A085098	001	
INWOOD LABS	25MG	A084776	001	
	25MG	A085067	001	
	50MG	A084776	002	
IPCA LABS LTD	12.5MG	A040807	001	Jul 20, 2007
	25MG	A040807	002	Jul 20, 2007
	50MG	A040807	003	Jul 20, 2007
IVAX SUB TEVA PHARMS	50MG	A084658	001	
+	100MG	A085022	001	
JUBILANT CADISTA	25MG	A040809	001	Sep 04, 2007
	50MG	A040809	002	Sep 04, 2007
LANNETT CO INC	25MG	A084325	001	
	50MG	A084324	001	
MAST MM	25MG	A086192	001	
	50MG	A086192	002	
MYLAN	25MG	A084880	001	
	50MG	A085112	001	
NORVIUM BIOSCIENCE	12.5MG	A040770	001	Jan 23, 2007
	25MG	A040735	002	Jan 23, 2007
	50MG	A040735	003	Jan 23, 2007

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

HYDROCHLOROTHIAZIDE

TABLET; ORAL

HYDROCHLOROTHIAZIDE

PRINSTON INC	25MG	A040412 001	Mar 29, 2002
	50MG	A040412 002	Mar 29, 2002
PVT FORM	50MG	A086597 001	
ROXANE	25MG	A085004 001	
	50MG	A084536 002	
	50MG	A085005 001	
SOLVAY	25MG	A085323 001	
SUN PHARM INDS INC	12.5MG	A040857 001	May 30, 2008
	25MG	A040810 001	Mar 27, 2007
	50MG	A040810 002	Mar 27, 2007
SUN PHARM INDUSTRIES	25MG	A083972 001	
	50MG	A083972 002	
	100MG	A083972 003	
SUPERPHARM	25MG	A088827 001	Dec 28, 1984
	50MG	A088828 001	Dec 28, 1984
	100MG	A088829 001	Dec 28, 1984
TEVA	25MG	A088924 001	Feb 07, 1985
	50MG	A088923 001	Feb 07, 1985
USL PHARMA	25MG	A087827 001	Apr 19, 1982
	50MG	A087752 001	Apr 19, 1982
VANGARD	25MG	A087638 001	
	50MG	A087610 001	
WARNER CHILCOTT	25MG	A087586 001	May 03, 1982
	50MG	A087587 001	May 03, 1982
WATSON LABS	25MG	A081189 001	Jan 24, 1992
	25MG	A083458 001	
	25MG	A085232 002	
	50MG	A083456 001	
	50MG	A085233 001	
	50MG	A086087 001	
	50MG	A086594 001	
	100MG	A081190 001	Jan 24, 1992
	100MG	A085099 001	
	100MG	A087002 001	
WATSON LABS TEVA	50MG	A083232 001	
WEST WARD	25MG	A084899 001	
WHITEWORTH TOWN PLSN	25MG	A083809 002	
	50MG	A083809 001	
	100MG	A085347 001	
HYDRODIURIL			
+ MERCK	25MG **	N011835 003	
+	50MG **	N011835 006	
+	100MG **	N011835 007	
ORETIC			
ABBVIE	25MG	N011971 001	
	50MG	N011971 002	
ZIDE			
SOLVAY	50MG	A083925 001	

HYDROCHLOROTHIAZIDE; IRBESARTAN

TABLET; ORAL

AVALIDE

+ SANOFI AVENTIS US	12.5MG; 75MG **	N020758 001	Sep 30, 1997
+	25MG; 300MG **	N020758 004	Mar 15, 2005

IRBESARTAN AND HYDROCHLOROTHIAZIDE

APOTEX INC	12.5MG; 150MG	A201505 001	Oct 15, 2012
	12.5MG; 300MG	A201505 002	Oct 15, 2012
ATLAS PHARMS LLC	12.5MG; 150MG	A203036 001	Jan 15, 2016
	12.5MG; 300MG	A203036 002	Jan 15, 2016
	25MG; 300MG	A203036 003	Jan 15, 2016
NORVIUM BIOSCIENCE	12.5MG; 150MG	A077969 001	Sep 27, 2012
	12.5MG; 300MG	A077969 002	Sep 27, 2012
	25MG; 300MG	A077969 003	Jul 20, 2016
TEVA	25MG; 300MG	A077369 003	Mar 30, 2012
UNICHEM	12.5MG; 150MG	A207018 001	Sep 19, 2017
	12.5MG; 300MG	A207018 002	Sep 19, 2017
WATSON LABS INC	12.5MG; 150MG	A091539 001	Oct 22, 2012

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

HYDROCHLOROTHIAZIDE; IRBESARTAN

TABLET; ORAL

IRBESARTAN AND HYDROCHLOROTHIAZIDE

12.5MG; 300MG

A091539 002 Oct 22, 2012

HYDROCHLOROTHIAZIDE; LABETALOL HYDROCHLORIDE

TABLET; ORAL

NORMOZIDE

SCHERING

25MG; 100MG

N019046 001 Apr 06, 1987

25MG; 200MG

N019046 002 Apr 06, 1987

25MG; 300MG

N019046 003 Apr 06, 1987

25MG; 400MG

N019046 004 Apr 06, 1987

TRANDATE HCT

GLAXOSMITHKLINE

25MG; 100MG

N019174 001 Apr 10, 1987

25MG; 200MG

N019174 002 Apr 10, 1987

25MG; 300MG

N019174 003 Apr 10, 1987

25MG; 400MG

N019174 004 Apr 10, 1987

HYDROCHLOROTHIAZIDE; LISINAPRIL

TABLET; ORAL

LISINAPRIL AND HYDROCHLOROTHIAZIDE

CHARTWELL RX

12.5MG; 10MG

A075776 001 Jul 01, 2002

12.5MG; 20MG

A075776 002 Jul 01, 2002

25MG; 20MG

A075776 003 Jul 01, 2002

EPIC PHARMA LLC

12.5MG; 10MG

A075926 001 Jul 01, 2002

12.5MG; 20MG

A075926 002 Jul 01, 2002

25MG; 20MG

A075926 003 Jul 01, 2002

HIKMA INTL PHARMS

12.5MG; 10MG

A076265 001 Jul 08, 2002

12.5MG; 20MG

A076265 002 Jul 08, 2002

25MG; 20MG

A076265 003 Jul 08, 2002

NORVIUM BIOSCIENCE

12.5MG; 10MG

A076113 001 Jul 01, 2002

12.5MG; 20MG

A076113 002 Jul 01, 2002

25MG; 20MG

A076113 003 Jul 01, 2002

TEVA

12.5MG; 10MG

A075869 001 Jul 01, 2002

12.5MG; 20MG

A075869 002 Jul 01, 2002

25MG; 20MG

A075869 003 Jul 01, 2002

PRINZIDE

+ MERCK

12.5MG; 10MG **

N019778 003 Nov 18, 1993

+

12.5MG; 20MG **

N019778 001 Feb 16, 1989

+

25MG; 20MG **

N019778 002 Feb 16, 1989

HYDROCHLOROTHIAZIDE; LOSARTAN POTASSIUM

TABLET; ORAL

LOSARTAN POTASSIUM AND HYDROCHLOROTHIAZIDE

APOTEX

12.5MG; 50MG

A090150 001 Oct 06, 2010

12.5MG; 100MG

A090150 002 Aug 11, 2010

25MG; 100MG

A090150 003 Oct 06, 2010

GRANULES

12.5MG; 50MG

A218015 001 Sep 29, 2023

12.5MG; 100MG

A218015 002 Sep 29, 2023

25MG; 100MG

A218015 003 Sep 29, 2023

HIKMA

12.5MG; 50MG

A077732 002 Oct 06, 2010

12.5MG; 100MG

A077732 001 Apr 06, 2010

25MG; 100MG

A077732 003 Oct 06, 2010

IPCA LABS LTD

12.5MG; 50MG

A201682 001 Mar 01, 2013

12.5MG; 100MG

A201682 002 Mar 01, 2013

25MG; 100MG

A201682 003 Mar 01, 2013

MYLAN

12.5MG; 50MG

A091652 001 Oct 06, 2010

12.5MG; 100MG

A091652 002 Apr 06, 2010

25MG; 100MG

A091652 003 Oct 06, 2010

TORRENT PHARMS

12.5MG; 50MG

A090528 001 Oct 06, 2010

12.5MG; 100MG

A090528 003 Apr 06, 2010

25MG; 100MG

A090528 002 Oct 06, 2010

WATSON LABS

12.5MG; 50MG

A200180 001 Jan 12, 2011

12.5MG; 100MG

A200180 002 Jan 12, 2011

25MG; 100MG

A200180 003 Jan 12, 2011

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

HYDROCHLOROTHIAZIDE; METHYLDOPA

TABLET; ORAL

ALDORIL 15

MERCK 15MG;250MG N013402 001

ALDORIL 25

MERCK 25MG;250MG N013402 002

ALDORIL D30

MERCK 30MG;500MG N013402 003

ALDORIL D50

MERCK 50MG;500MG N013402 004

METHYLDOPA AND HYDROCHLOROTHIAZIDE

CHARTWELL RX 15MG;250MG A070182 001 Jan 15, 1986

25MG;250MG A070183 001 Jan 15, 1986

30MG;500MG A070543 001 Jan 15, 1986

50MG;500MG A070544 001 Jan 15, 1986

DAVA PHARMS INC 15MG;250MG A072507 001 Jun 02, 1989

25MG;250MG A072508 001 Jun 02, 1989

30MG;500MG A072509 001 Jun 02, 1989

50MG;500MG A072510 001 Jun 02, 1989

IVAX SUB TEVA PHARMS 15MG;250MG A071458 001 Mar 08, 1988

25MG;250MG A071459 001 Mar 08, 1988

30MG;500MG A071460 001 Mar 08, 1988

50MG;500MG A071461 001 Mar 08, 1988

PARKE DAVIS 15MG;250MG A071897 001 Nov 23, 1987

25MG;250MG A071898 001 Nov 23, 1987

30MG;500MG A071899 001 Nov 23, 1987

50MG;500MG A071900 001 Nov 23, 1987

PUREPAC PHARM 15MG;250MG A070853 001 Oct 08, 1986

25MG;250MG A070688 001 Apr 24, 1986

30MG;500MG A070854 001 Oct 08, 1986

50MG;500MG A070689 001 Apr 24, 1986

RISING 15MG;250MG A070265 002 Jan 23, 1986

25MG;250MG A070265 001 Jan 23, 1986

SANDOZ 15MG;250MG A070829 001 Mar 09, 1987

25MG;250MG A070830 001 Mar 09, 1987

STRIDES PHARMA 15MG;250MG A070616 001 Feb 02, 1987

25MG;250MG A070612 001 Feb 02, 1987

30MG;500MG A070613 001 Feb 02, 1987

50MG;500MG A070614 001 Feb 02, 1987

TEVA 15MG;250MG A071819 001 Apr 08, 1988

25MG;250MG A071820 001 Apr 08, 1988

30MG;500MG A071821 001 Apr 08, 1988

50MG;500MG A071822 001 Apr 08, 1988

WATSON LABS 15MG;250MG A070365 001 Mar 19, 1986

15MG;250MG A070958 001 Feb 06, 1989

15MG;250MG A071920 001 Aug 29, 1988

25MG;250MG A070366 001 Apr 16, 1986

25MG;250MG A070959 001 Jan 19, 1989

25MG;250MG A071921 001 Aug 29, 1988

30MG;500MG A070367 001 Mar 19, 1986

30MG;500MG A071069 001 Jan 19, 1989

30MG;500MG A071922 001 Aug 29, 1988

50MG;500MG A070368 001 Apr 16, 1986

50MG;500MG A070960 001 Feb 06, 1989

50MG;500MG A071923 001 Aug 29, 1988

HYDROCHLOROTHIAZIDE; METOPROLOL SUCCINATE

TABLET, EXTENDED RELEASE; ORAL

DUTOPROL

+ CONCORDIA 12.5MG;EQ 25MG TARTRATE ** N021956 001 Aug 28, 2006

+ 12.5MG;EQ 50MG TARTRATE ** N021956 002 Aug 28, 2006

+ 12.5MG;EQ 100MG TARTRATE ** N021956 003 Aug 28, 2006

HYDROCHLOROTHIAZIDE; METOPROLOL TARTRATE

TABLET; ORAL

LOPRESSOR HCT

+ VALIDUS PHARMS 25MG;100MG ** N018303 002 Dec 31, 1984

+ 50MG;100MG ** N018303 003 Dec 31, 1984

METOPROLOL TARTRATE AND HYDROCHLOROTHIAZIDE

SUN PHARM INDS 25MG;50MG A090654 001 Jan 19, 2012

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

HYDROCHLOROTHIAZIDE; METOPROLOL TARTRATE

TABLET; ORAL

METOPROLOL TARTRATE AND HYDROCHLOROTHIAZIDE

25MG;100MG

A090654 002 Jan 19, 2012

50MG;100MG

A090654 003 Jan 19, 2012

HYDROCHLOROTHIAZIDE; MOEXIPRIL HYDROCHLORIDE

TABLET; ORAL

MOEXIPRIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE

CHARTWELL RX

12.5MG; 7.5MG

A090096 001 Sep 25, 2008

12.5MG; 15MG

A090096 002 Sep 25, 2008

25MG; 15MG

A090096 003 Sep 25, 2008

HERITAGE

12.5MG; 7.5MG

A202150 001 Mar 07, 2014

12.5MG; 15MG

A202150 002 Mar 07, 2014

25MG; 15MG

A202150 003 Mar 07, 2014

UNIRETIC

UCB INC

12.5MG; 7.5MG **

N020729 001 Jun 27, 1997

12.5MG; 15MG **

N020729 003 Feb 14, 2002

25MG; 15MG **

N020729 002 Jun 27, 1997

HYDROCHLOROTHIAZIDE; OLMESARTAN MEDOXOMIL

TABLET; ORAL

OLMESARTAN MEDOXOMIL AND HYDROCHLOROTHIAZIDE

NATCO PHARMA USA

12.5MG; 20MG

A078827 001 Oct 26, 2016

12.5MG; 40MG

A078827 002 Oct 26, 2016

25MG; 40MG

A078827 003 Oct 26, 2016

TEVA PHARMS USA

12.5MG; 20MG

A200532 001 Apr 24, 2017

12.5MG; 40MG

A200532 002 Apr 24, 2017

25MG; 40MG

A200532 003 Apr 24, 2017

TORRENT

12.5MG; 20MG

A206515 001 May 03, 2017

12.5MG; 40MG

A206515 002 May 03, 2017

25MG; 40MG

A206515 003 May 03, 2017

ZYDUS PHARMS

12.5MG; 20MG

A206377 001 Feb 24, 2023

12.5MG; 40MG

A206377 002 Feb 24, 2023

25MG; 40MG

A206377 003 Feb 24, 2023

HYDROCHLOROTHIAZIDE; PINDOLOL

TABLET; ORAL

VISKAZIDE

NOVARTIS

25MG; 5MG

N018872 001 Jul 22, 1987

25MG; 10MG

N018872 002 Jul 22, 1987

HYDROCHLOROTHIAZIDE; PROPRANOLOL HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

INDERIDE LA 120/50

WYETH AYERST

50MG; 120MG

N019059 002 Jul 03, 1985

INDERIDE LA 160/50

WYETH AYERST

50MG; 160MG

N019059 003 Jul 03, 1985

INDERIDE LA 80/50

WYETH AYERST

50MG; 80MG

N019059 001 Jul 03, 1985

TABLET; ORAL

INDERIDE-40/25

+ WYETH PHARMS INC

25MG; 40MG **

N018031 001

INDERIDE-80/25

+ WYETH PHARMS INC

25MG; 80MG **

N018031 002

PROPRANOLOL HYDROCHLORIDE & HYDROCHLOROTHIAZIDE

DURAMED PHARMS BARR

25MG; 40MG

A071126 001 Mar 02, 1987

25MG; 80MG

A071127 001 Mar 02, 1987

PROPRANOLOL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE

ACTAVIS ELIZABETH

25MG; 40MG

A070851 001 May 15, 1986

25MG; 80MG

A070852 001 May 15, 1986

ANI PHARMS

25MG; 40MG

A070705 002 Oct 01, 1986

25MG; 40MG

A072043 002 Mar 14, 1988

25MG; 80MG

A070705 001 Oct 01, 1986

25MG; 80MG

A072043 001 Mar 14, 1988

CHARTWELL RX

25MG; 40MG

A071060 001 Aug 26, 1987

25MG; 80MG

A071061 001 Aug 26, 1987

IVAX SUB TEVA PHARMS

25MG; 40MG

A071552 001 Dec 01, 1988

25MG; 80MG

A071553 001 Dec 01, 1988

RISING

25MG; 40MG

A070947 002 Mar 04, 1987

25MG; 80MG

A070947 001 Apr 01, 1987

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

HYDROCHLOROTHIAZIDE; PROPRANOLOL HYDROCHLORIDE

TABLET; ORAL

PROPRANOLOL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE

WARNER CHILCOTT	25MG; 40MG	A071771	001	Jan 26, 1988
	25MG; 80MG	A071772	001	Jan 26, 1988
WATSON LABS	25MG; 40MG	A070301	001	Apr 18, 1986
	25MG; 40MG	A071498	001	Dec 18, 1991
	25MG; 80MG	A070305	001	Apr 18, 1986
	25MG; 80MG	A071501	001	Dec 18, 1991

HYDROCHLOROTHIAZIDE; QUINAPRIL HYDROCHLORIDE

TABLET; ORAL

ACCURETIC

+	PFIZER PHARMS	12.5MG; EQ 10MG BASE **	N020125	001	Dec 28, 1999
+		12.5MG; EQ 20MG BASE **	N020125	002	Dec 28, 1999
+		25MG; EQ 20MG BASE **	N020125	003	Dec 28, 1999

QUINAPRIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE

INVAGEN PHARMS	12.5MG; EQ 10MG BASE	A201356	001	Apr 20, 2011
	12.5MG; EQ 20MG BASE	A201356	002	Apr 20, 2011
	25MG; EQ 20MG BASE	A201356	003	Apr 20, 2011
NORVIUM BIOSCIENCE	12.5MG; EQ 10MG BASE	A077093	001	Mar 28, 2005
	12.5MG; EQ 20MG BASE	A077093	002	Mar 28, 2005
	25MG; EQ 20MG BASE	A077093	003	Mar 28, 2005
SUN PHARM INDS LTD	12.5MG; EQ 10MG BASE	A078211	001	Mar 04, 2009
	12.5MG; EQ 20MG BASE	A078211	002	Mar 04, 2009
	25MG; EQ 20MG BASE	A078211	003	Mar 04, 2009

HYDROCHLOROTHIAZIDE; RESERPINE

TABLET; ORAL

H.R.-50

WHITEWORTH TOWN PLSN	50MG; 0.125MG	A085338	001	
----------------------	---------------	---------	-----	--

HYDRO-RESERP

ABC HOLDING	50MG; 0.125MG	A084714	002	Jun 29, 1982
-------------	---------------	---------	-----	--------------

HYDRO-SERP "25"

SANDOZ	25MG; 0.125MG	A084827	001	
--------	---------------	---------	-----	--

HYDRO-SERP "50"

SANDOZ	50MG; 0.125MG	A085213	001	
--------	---------------	---------	-----	--

HYDROCHLOROTHIAZIDE W/ RESERPINE

IVAX SUB TEVA PHARMS	25MG; 0.1MG	A083572	001	
	25MG; 0.125MG	A083571	001	
	50MG; 0.1MG	A083568	001	
	50MG; 0.125MG	A083573	001	
PHARMERAL	25MG; 0.125MG	A085421	001	
	50MG; 0.125MG	A085420	001	
ROXANE	50MG; 0.125MG	A084603	001	
WATSON LABS	25MG; 0.125MG	A084466	001	
	25MG; 0.125MG	A085317	001	
	25MG; 0.125MG	A086330	002	
	50MG; 0.125MG	A083666	001	
	50MG; 0.125MG	A084467	001	
	50MG; 0.125MG	A086331	001	

HYDROPRES 25

MERCK	25MG; 0.125MG	N011958	002	
-------	---------------	---------	-----	--

HYDROPRES 50

MERCK	50MG; 0.125MG	N011958	003	
-------	---------------	---------	-----	--

RESERPINE AND HYDROCHLOROTHIAZIDE

BARR	25MG; 0.125MG	A084580	001	
	50MG; 0.125MG	A084579	001	
SANDOZ	50MG; 0.125MG	A088200	001	Jan 31, 1984

RESERPINE AND HYDROCHLOROTHIAZIDE-50

WEST WARD	50MG; 0.125MG	A088189	001	May 10, 1984
-----------	---------------	---------	-----	--------------

SERPASIL-ESIDRIX #1

NOVARTIS	25MG; 0.1MG	N011878	003	
----------	-------------	---------	-----	--

SERPASIL-ESIDRIX #2

NOVARTIS	50MG; 0.1MG	N011878	005	
----------	-------------	---------	-----	--

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

HYDROCHLOROTHIAZIDE; SPIRONOLACTONE

TABLET; ORAL

ALDACTAZIDE

+ PFIZER 50MG; 50MG N012616 005 Dec 30, 1982

SPIRONOLACTONE AND HYDROCHLOROTHIAZIDE

ASCOT 25MG; 25MG A088025 001 Nov 23, 1984

CHARTWELL RX 25MG; 25MG A086881 001

MUTUAL PHARM 25MG; 25MG A087267 001

PUREPAC PHARM 25MG; 25MG A087999 001 Nov 06, 1985

SUPERPHARM 25MG; 25MG A089137 001 Aug 26, 1985

WATSON LABS 25MG; 25MG A087398 001

SPIRONOLACTONE W/ HYDROCHLOROTHIAZIDE

IVAX PHARMS 25MG; 25MG A087004 002 May 24, 1982

LEDERLE 25MG; 25MG A087511 001

PARKE DAVIS 25MG; 25MG A087948 001 Feb 22, 1983

PUREPAC PHARM 25MG; 25MG A088054 001 Aug 18, 1983

UPSHER SMITH 25MG; 25MG A087553 001

USL PHARMA 25MG; 25MG A087651 001

VANGARD 25MG; 25MG A087655 001

WATSON LABS 25MG; 25MG A085974 001

25MG; 25MG A086026 001

HYDROCHLOROTHIAZIDE; TELMISARTAN

TABLET; ORAL

TELMISARTAN AND HYDROCHLOROTHIAZIDE

MACLEODS PHARMS LTD 12.5MG; 40MG A204169 001 Nov 02, 2015

12.5MG; 80MG A204169 002 Nov 02, 2015

25MG; 80MG A204169 003 Nov 02, 2015

NATCO 12.5MG; 40MG A091648 001 Feb 25, 2014

12.5MG; 80MG A091648 002 Feb 25, 2014

25MG; 80MG A091648 003 Feb 25, 2014

TORRENT 12.5MG; 40MG A201192 001 Feb 25, 2014

12.5MG; 80MG A201192 002 Feb 25, 2014

25MG; 80MG A201192 003 Feb 25, 2014

HYDROCHLOROTHIAZIDE; TIMOLOL MALEATE

TABLET; ORAL

TIMOLIDE 10-25

MERCK 25MG; 10MG N018061 001

HYDROCHLOROTHIAZIDE; TRIAMTERENE

CAPSULE; ORAL

DYAZIDE

+ GLAXOSMITHKLINE LLC 25MG; 37.5MG ** N016042 003 Mar 03, 1994

25MG; 50MG ** N016042 002

TRIAMTERENE AND HYDROCHLOROTHIAZIDE

ANI PHARMS 25MG; 37.5MG A074970 001 Jan 06, 1998

25MG; 50MG A074259 001 Mar 30, 1995

CHARTWELL RX 25MG; 50MG A073191 001 Jul 31, 1991

DURAMED PHARMS BARR 25MG; 37.5MG A075052 001 Jun 18, 1999

NORVIUM BIOSCIENCE 25MG; 37.5MG A074701 001 Jun 07, 1996

NOVARTIS 25MG; 37.5MG A074857 001 Sep 09, 1997

VITARINE 25MG; 50MG A071737 001 Feb 12, 1988

TABLET; ORAL

TRIAMTERENE AND HYDROCHLOROTHIAZIDE

AM THERAP 50MG; 75MG A072022 001 Apr 17, 1988

ANI PHARMS 50MG; 75MG A073467 001 Jan 31, 1996

PLIVA 25MG; 37.5MG A074026 001 Apr 26, 1996

QUANTUM PHARMICS 50MG; 75MG A071980 001 Apr 17, 1988

WATSON LABS 50MG; 75MG A071969 001 Apr 17, 1988

HYDROCHLOROTHIAZIDE; VALSARTAN

TABLET; ORAL

VALSARTAN AND HYDROCHLOROTHIAZIDE

APOTEX INC 12.5MG; 80MG A203026 001 Mar 21, 2013

12.5MG; 160MG A203026 002 Mar 21, 2013

12.5MG; 320MG A203026 003 Mar 21, 2013

25MG; 160MG A203026 004 Mar 21, 2013

25MG; 320MG A203026 005 Mar 21, 2013

WATSON LABS TEVA 12.5MG; 80MG A091519 001 Mar 21, 2013

12.5MG; 160MG A091519 002 Mar 21, 2013

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

HYDROCHLOROTHIAZIDE; VALSARTAN

TABLET; ORAL

VALSARTAN AND HYDROCHLOROTHIAZIDE

	12.5MG; 320MG	A091519 003	Mar 21, 2013
	25MG; 160MG	A091519 004	Mar 21, 2013
	25MG; 320MG	A091519 005	Mar 21, 2013
ZYDUS LIFESCIENCES	12.5MG; 80MG	A203000 001	Mar 15, 2019
	12.5MG; 160MG	A203000 002	Mar 15, 2019
	12.5MG; 320MG	A203000 003	Mar 15, 2019
	25MG; 160MG	A203000 004	Mar 15, 2019
	25MG; 320MG	A203000 005	Mar 15, 2019

HYDROCODONE BITARTRATE

CAPSULE, EXTENDED RELEASE; ORAL

ZOHYDRO ER

+	RECRO GAINESVILLE	10MG	N202880 001	Oct 25, 2013
+		15MG	N202880 002	Oct 25, 2013
+		20MG	N202880 003	Oct 25, 2013
+		30MG	N202880 004	Oct 25, 2013
+		40MG	N202880 005	Oct 25, 2013
+		50MG	N202880 006	Oct 25, 2013

TABLET, EXTENDED RELEASE; ORAL

HYSINGLA ER

+	PURDUE PHARMA LP	120MG	N206627 007	Nov 20, 2014
---	------------------	-------	-------------	--------------

VANTRELA ER

+	TEVA BRANDED PHARM	15MG	N207975 001	Jan 17, 2017
+		30MG	N207975 002	Jan 17, 2017
+		45MG	N207975 003	Jan 17, 2017
+		60MG	N207975 004	Jan 17, 2017
+		90MG	N207975 005	Jan 17, 2017

HYDROCODONE BITARTRATE; IBUPROFEN

TABLET; ORAL

HYDROCODONE

	NOSTRUM LABS INC	5MG; 200MG	A077723 003	Nov 06, 2006
--	------------------	------------	-------------	--------------

HYDROCODONE BITARTRATE AND IBUPROFEN

	ANI PHARMS	5MG; 200MG	A077454 001	Jun 23, 2010
	NOSTRUM LABS INC	7.5MG; 200MG	A077723 001	Nov 06, 2006
		10MG; 200MG	A077723 002	Nov 06, 2006
	SUN PHARM INDS INC	2.5MG; 200MG	A091633 001	May 28, 2013
		5MG; 200MG	A091633 002	May 28, 2013
		7.5MG; 200MG	A091633 003	May 28, 2013
		10MG; 200MG	A091633 004	May 28, 2013
	TEVA	7.5MG; 200MG	A076023 001	Apr 11, 2003

REPREXAIN

	AMNEAL PHARMS NY	2.5MG; 200MG	A076642 003	Oct 19, 2007
		10MG; 200MG	A076642 004	Oct 19, 2007

VICOPROFEN

+	ABBVIE	7.5MG; 200MG **	N020716 001	Sep 23, 1997
---	--------	-----------------	-------------	--------------

HYDROCODONE BITARTRATE; PHENYLPROPANOLAMINE HYDROCHLORIDE

SYRUP; ORAL

CODAMINE

	ALPHARMA US PHARMS	5MG/5ML; 25MG/5ML	A075103 001	Sep 29, 2000
--	--------------------	-------------------	-------------	--------------

HYDROCODONE BITARTRATE; PSEUDOEPHEDRINE HYDROCHLORIDE

SOLUTION; ORAL

HYDROCODONE BITARTRATE AND PSEUDOEPHEDRINE HYDROCHLORIDE

	MAYNE PHARMA INC	5MG/5ML; 60MG/5ML	A205658 001	Nov 17, 2015
	PADAGIS US	5MG/5ML; 60MG/5ML	A204658 001	Apr 29, 2014
	SCIEGEN PHARMS INC	5MG/5ML; 60MG/5ML	A206661 001	Jan 23, 2019
	TRIS PHARMA INC	5MG/5ML; 60MG/5ML	A203839 001	Oct 28, 2014

REZIRA

+	PERSION	5MG/5ML; 60MG/5ML **	N022442 001	Jun 08, 2011
---	---------	----------------------	-------------	--------------

HYDROCORTAMATE HYDROCHLORIDE

OINTMENT; TOPICAL

MAGNACORT

	PFIZER	0.5%	N010554 001	
--	--------	------	-------------	--

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

HYDROCORTISONE

AEROSOL; TOPICAL

AEROSEB-HC

ALLERGAN HERBERT 0.5% A085805 001

CREAM; TOPICAL

CORT-DOME

BAYER PHARMS 0.5% N009585 003

1% N009585 001

DERMACORT

MONARCH PHARMS 1% A083011 002

ELDECORT

VALEANT PHARM INTL 1% A080459 001

2.5% A084055 001

FLEXICORT

WESTWOOD SQUIBB 0.5% A087136 003 Apr 08, 1982

1% A087136 002 Apr 08, 1982

2.5% A087136 001 Apr 08, 1982

H-CORT

PHARM ASSOC 0.5% A086823 001

HC #1

BAYER PHARMS 0.5% A080438 001

HC #4

BAYER PHARMS 1% A080438 002

HC (HYDROCORTISONE)

C AND M PHARMA 0.5% A080482 003

1% A080482 004

HI-COR

C AND M PHARMA 2.5% A080483 001

HYDROCORTISONE

ALPHARMA US PHARMS 2.5% A089754 001 Feb 01, 1989

ALTANA 0.5% A080848 002

1% A080848 003

AMBIX 1% A086080 001

2.5% A086271 001

CHARTWELL MOLECULAR 2.5% A040503 001 Mar 12, 2004

ENCUBE 2.5% A203810 001 Jul 23, 2018

EVERYLIFE 0.5% A080452 001

1% A080452 002

G AND W LABS 1% A084059 001

INGRAM PHARM 0.5% A080456 002

1% A080456 003

IVAX PHARMS 1% A085733 001

NASKA 1% A089706 001 Mar 10, 1988

PERRIGO NEW YORK 0.5% A084970 002

1% A085026 001

PHARMADERM 1% A088845 001 Feb 27, 1986

2.5% A089413 001 Dec 16, 1986

PHARMAFAIR 1% A087838 001 Jul 28, 1982

STIEFEL 1% A086170 001

SYOSSET 0.5% A085527 001

TARO 0.5% A086154 001

1% A086155 001

TEVA 0.5% A080400 002

1% A080400 003

1% A085191 001

2.5% A080400 004

TOPIDERM 1% A089273 001 Feb 17, 1989

USL PHARMA 1% A088027 001 Sep 27, 1983

2.5% A088029 001 Sep 27, 1983

WHITEWORTH TOWN PLSN 1% A080496 002

HYTONE

+ VALEANT INTL 1% ** A080472 003

+ 2.5% ** A080472 004

NOGENIC HC

IVAX PHARMS 1% A087427 001 Apr 04, 1988

NUTRACORT

BAUSCH 0.5% A080442 002

1% A080442 003

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

HYDROCORTISONE

CREAM; TOPICAL

PENECORT

ALLERGAN HERBERT 1% A088216 001 Jun 06, 1984

PROCTOCORT

MONARCH PHARMS 1% A083011 001

SYNACORT

BAUSCH 0.5% A087459 001

+ 1% A087458 001

+ 2.5% A087457 001

ENEMA; RECTAL

HYDROCORTISONE

TEVA PHARMS 100MG/60ML A074171 001 May 27, 1994

GEL; TOPICAL

NUTRACORT

HEALTHPOINT 1% A084698 001

PENECORT

ALLERGAN HERBERT 1% A088215 001 Jun 06, 1984

INJECTABLE; INJECTION

CORTEF

PHARMACIA AND UPJOHN 50MG/ML N009864 001

LOTION; TOPICAL

ACTICORT

BAKER NORTON 1% A086535 001

ALA-CORT

CROWN LABS 1% A083201 001

BALNEOL-HC

SOLVAY 1% A088041 001 Dec 03, 1982

BETA-HC

BETA DERMAC 1% A089495 001 Jan 25, 1988

CETACORT

BAUSCH 0.5% A080426 002

1% A080426 001

CORT-DOME

BAYER PHARMS 0.5% N009895 003

1% N009895 001

DERMACORT

SOLVAY 0.5% A084573 002

1% A086462 001

EPICORT

BLULINE 0.5% A083219 002

GLYCORT

HERAN 1% A087489 001 Oct 03, 1983

H-CORT

PHARM ASSOC 0.5% A086824 001

HYDROCORTISONE

ALPHARMA US PHARMS 0.5% A087317 001 Jun 07, 1982

1% A087315 001 Jun 07, 1982

CHARTWELL MOLECULAR 2.5% A040417 001 Jul 30, 2003

ENCUBE 2.5% A203804 001 Jul 27, 2018

FOUGERA PHARMS 2.5% A040351 001 Jul 25, 2000

MERICON 0.5% A085282 001

1% A085282 002 Feb 26, 1987

NASKA 1% A089705 001 Apr 25, 1988

PERRIGO NEW YORK 0.5% A085662 001

1% A085663 001

TARO 1% A089024 001 Feb 12, 1986

HYTONE

+ VALEANT INTL 1% ** A080473 003

+ 2.5% ** A080473 004 Nov 30, 1982

NUTRACORT

DOW PHARM 0.5% A080443 002

1% A080443 003

2.5% A087644 001 Aug 24, 1982

STIE-CORT

PADAGIS US 1% A089066 001 Nov 25, 1985

OINTMENT; TOPICAL

CORTRIL

PFIZER GLOBAL 1% N009176 001

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

HYDROCORTISONE

OINTMENT;TOPICAL

CORTRIL	2.5%		N009176 002	
HC (HYDROCORTISONE)				
C AND M PHARMA	0.5%		A080481 001	
	1%		A080481 002	
HYDROCORTISONE				
ACTAVIS MID ATLANTIC	1%		A087796 001	Oct 13, 1982
ALTANA	0.5%		A080489 002	
	1%		A080489 003	
AMBIX	1%		A086079 001	
	2.5%		A086272 001	
NASKA	1%		A089704 001	Mar 10, 1988
PERRIGO NEW YORK	0.5%		A084969 003	
	1%		A085028 001	
PHARMADERM	1%		A088842 001	Feb 09, 1987
TARO	0.5%		A086256 001	
	2.5%		A040310 001	Dec 29, 2000
USL PHARMA	1%		A088061 001	Sep 27, 1983
	2.5%		A088039 001	Sep 27, 1983
HYTONE				
+ DERMIK LABS	1% **		A080474 003	
+	2.5% **		A080474 004	
PENECORT				
ALLERGAN HERBERT	2.5%		A088217 001	Jun 06, 1984
POWDER;FOR RX COMPOUNDING				
H-CORT				
TORCH	100%		A087834 001	Mar 29, 1982
HYDRO-RX				
X GEN PHARMS	100%		A085982 001	
HYDROCORTISONE				
PADDOCK LLC	100%		A088082 001	Apr 08, 1983
SOLUTION;TOPICAL				
PENECORT				
+ ALLERGAN HERBERT	1%		A088214 001	Jun 06, 1984
TEXACORT				
MISSION PHARMA	1%		A080425 001	
TABLET;ORAL				
CORTRIL				
PFIZER	10MG		N009127 005	
	20MG		N009127 003	
HYDROCORTISONE				
BARR	20MG		A083999 001	
CHARTWELL MOLECULAR	20MG		A085070 001	
ELKINS SINN	20MG		A080624 001	
FERRANTE	10MG		A080568 001	
	20MG		A080568 002	
HIKMA INTL PHARMS	5MG		A083365 002	Feb 23, 2015
	10MG		A083365 003	Feb 23, 2015
	20MG		A083365 001	
IMPAX LABS	20MG		A080781 001	
INWOOD LABS	20MG		A080732 001	
NEXGEN PHARMA INC	20MG		A083140 001	
PANRAY	10MG		N009659 001	
	20MG		N009659 002	
PARKE DAVIS	20MG		A084243 001	
PUREPAC PHARM	10MG		A084247 003	Aug 31, 1982
	20MG		A080395 001	
	20MG		A084247 002	
ROXANE	10MG		A088539 001	Mar 21, 1984
SANDOZ	20MG		A080642 002	
STRIDES PHARMA	5MG		A040761 001	Jul 16, 2007
	10MG		A040761 002	Jul 16, 2007
	20MG		A040761 003	Jul 16, 2007
WATSON LABS	20MG		A080355 001	
WHITEWORTH TOWN PLSN	10MG		A080344 001	
	20MG		A080344 002	

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

HYDROCORTISONE

TABLET; ORAL

HYDROCORTONE

MERCK

10MG

N008506 007

20MG

N008506 011

TABLET; VAGINAL

CORTRIL

PFIPHARMECS

10MG

N009796 001

HYDROCORTISONE ACETATE

CREAM; TOPICAL

CARMOL HC

+ FOUGERA PHARMS

1%

A080505 001

HEMSOL-HC

ABLE

1%

A081274 001 Jun 19, 1992

HYDROCORTISONE ACETATE

CENCI

1%

A080419 001 Jan 25, 1982

IMPERIUM

2.5%

A040259 001 Jul 29, 1999

PARKE DAVIS

1%

A089914 001 Jan 03, 1989

PUREPAC PHARM

0.5%

A086050 001

+

1%

A086052 001

MICORT-HC

SEBELA IRELAND LTD

2%

A040398 001 Mar 29, 2002

U-CORT

TARO

1%

A089472 001 Jun 13, 1988

INJECTABLE; INJECTION

CORTEF ACETATE

PHARMACIA AND UPJOHN

50MG/ML

N009378 002

CORTRIL

PFIZER

25MG/ML

N009164 001

HYDROCORTISONE ACETATE

BEL MAR

25MG/ML

A083739 001

50MG/ML

A083739 002

EPIC PHARMA LLC

25MG/ML

N009637 001

50MG/ML

N009637 002

WATSON LABS

25MG/ML

A083128 001

25MG/ML

A083759 001

50MG/ML

A083759 002

50MG/ML

A085214 001

HYDROCORTONE

MERCK

25MG/ML

N008228 001

50MG/ML

N008228 004

LOTION; TOPICAL

DRICORT

INGRAM PHARM

0.5%

A086207 001

OINTMENT; OPHTHALMIC

HYDROCORTISONE ACETATE

FERA PHARMS

0.5%

A080828 001

OINTMENT; OPHTHALMIC, OTIC

HYDROCORTONE

MERCK

1.5%

N009018 003

OINTMENT; TOPICAL

CORTEF ACETATE

PHARMACIA AND UPJOHN

1%

N008917 002

+

2.5% **

N008917 001

PASTE; TOPICAL

ORABASE HCA

COLGATE

0.5%

A083205 001

POWDER; FOR RX COMPOUNDING

HYDROCORTISONE ACETATE

X GEN PHARMS

100%

A085981 001

HYDROCORTISONE ACETATE; NEOMYCIN SULFATE

CREAM; TOPICAL

NEO-CORTEF

PHARMACIA AND UPJOHN 1%;EQ 3.5MG BASE/GM

A061049 001

2.5%;EQ 3.5MG BASE/GM

A061049 002

OINTMENT; OPHTHALMIC

NEO-CORTEF

PHARMACIA AND UPJOHN 0.5%;EQ 3.5MG BASE/GM

A060610 001

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

HYDROCORTISONE ACETATE; NEOMYCIN SULFATE

OINTMENT;OPHTHALMIC

NEO-CORTEF

1.5%;EQ 3.5MG BASE/GM

A060610 002

OINTMENT;TOPICAL

NEO-CORTEF

PHARMACIA AND UPJOHN 0.5%;EQ 3.5MG BASE/GM

A060751 001

1%;EQ 3.5MG BASE/GM

A060751 002

2.5%;EQ 3.5MG BASE/GM

A060751 003

SUSPENSION/DROPS;OPHTHALMIC

COR-OTICIN

EPIC PHARMA LLC 1.5%;EQ 3.5MG BASE/ML

A060188 001

NEO-CORTEF

PHARMACIA AND UPJOHN 0.5%;EQ 3.5MG BASE/ML

A060612 002

1.5%;EQ 3.5MG BASE/ML

A060612 001

HYDROCORTISONE ACETATE; NEOMYCIN SULFATE; POLYMYXIN B SULFATE

CREAM;TOPICAL

CORTISPORIN

+ MONARCH PHARMS 0.5%;EQ 3.5MG BASE/GM;10,000 UNITS/GM

N050218 001 Aug 09, 1985

**

HYDROCORTISONE ACETATE; OXYTETRACYCLINE HYDROCHLORIDE

SUSPENSION;OPHTHALMIC

TERRA-CORTRIL

PFIZER 1.5%;EQ 5MG BASE/ML

A061016 001

HYDROCORTISONE ACETATE; PRAMOXINE HYDROCHLORIDE

AEROSOL, METERED;TOPICAL

HYDROCORTISONE ACETATE 1% AND PRAMOXINE HYDROCHLORIDE 1%

GENUS 1%;1%

A089440 001 May 17, 1988

LOTION;TOPICAL

PRAMOSONE

FERNDALE LABS 0.5%;1%

A083213 002

HYDROCORTISONE BUTYRATE

CREAM;TOPICAL

LOCOID

YAMANOUCHI 0.1%

N018795 001 Jan 07, 1983

LOTION;TOPICAL

LOCOID

+ BAUSCH 0.1%

N022076 001 May 18, 2007

OINTMENT;TOPICAL

LOCOID

YAMANOUCHI 0.1%

N019106 001 Jul 03, 1984

SOLUTION;TOPICAL

LOCOID

YAMANOUCHI 0.1%

N019819 001 Sep 15, 1988

HYDROCORTISONE CYPIONATE

SUSPENSION;ORAL

CORTEF

PHARMACIA AND UPJOHN EQ 10MG BASE/5ML

N009900 001

HYDROCORTISONE PROBUTATE

CREAM;TOPICAL

PANDEL

+ ANI PHARMS 0.1%

N020453 001 Feb 28, 1997

HYDROCORTISONE SODIUM PHOSPHATE

INJECTABLE;INJECTION

HYDROCORTONE

+ MERCK EQ 50MG BASE/ML **

N012052 001

HYDROCORTISONE SODIUM SUCCINATE

INJECTABLE;INJECTION

A-HYDROCORT

ABBOTT EQ 100MG BASE/VIAL

A085928 001

EQ 100MG BASE/VIAL

A089577 001 Apr 11, 1989

EQ 250MG BASE/VIAL

A089578 001 Apr 11, 1989

EQ 500MG BASE/VIAL

A089579 001 Apr 11, 1989

EQ 1GM BASE/VIAL

A089580 001 Apr 11, 1989

HOSPIRA EQ 100MG BASE/VIAL

A040666 001 Apr 06, 2006

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

HYDROCORTISONE SODIUM SUCCINATE

INJECTABLE; INJECTION

A-HYDROCORT

EQ 100MG BASE/VIAL	A085929	001
EQ 250MG BASE/VIAL	A085930	001
EQ 500MG BASE/VIAL	A085931	001
EQ 1GM BASE/VIAL	A085932	001

HYDROCORTISONE SODIUM SUCCINATE

ABRAXIS PHARM

EQ 100MG BASE/VIAL	A088667	001	Jun 08, 1984
EQ 100MG BASE/VIAL	A088712	001	Jun 08, 1984
EQ 250MG BASE/VIAL	A088668	001	Jun 08, 1984
EQ 500MG BASE/VIAL	A088669	001	Jun 08, 1984
EQ 1GM BASE/VIAL	A088670	001	Jun 08, 1984

BAXTER HLTHCARE

EQ 100MG BASE/VIAL	A086619	001
EQ 250MG BASE/VIAL	A087567	001
EQ 500MG BASE/VIAL	A087568	001
EQ 1GM BASE/VIAL	A087569	001

INTL MEDICATION

EQ 100MG BASE/VIAL	A087532	001	Mar 19, 1982
--------------------	---------	-----	--------------

WATSON LABS

EQ 100MG BASE/VIAL	A084737	002
EQ 100MG BASE/VIAL	A084738	001
EQ 250MG BASE/VIAL	A084737	001
EQ 500MG BASE/VIAL	A084747	001
EQ 1GM BASE/VIAL	A084748	001

HYDROCORTISONE VALERATE

CREAM; TOPICAL

HYDROCORTISONE VALERATE

COSETTE

0.2%	A074489	001	Aug 12, 1998
------	---------	-----	--------------

WESTCORT

+ SUN PHARM INDS INC

0.2% **	N017950	001
---------	---------	-----

OINTMENT; TOPICAL

HYDROCORTISONE VALERATE

FOUGERA PHARMS

0.2%	A075085	001	Jul 31, 2001
------	---------	-----	--------------

WESTCORT

+ SUN PHARM INDS INC

0.2% **	N018726	001	Aug 08, 1983
---------	---------	-----	--------------

HYDROCORTISONE; NEOMYCIN SULFATE

CREAM; TOPICAL

NEO-CORT-DOME

BAYER PHARMS

0.5%;EQ 3.5MG BASE/GM	N050237	006	Jun 05, 1984
1%;EQ 3.5MG BASE/GM	N050237	005	Jun 05, 1984

HYDROCORTISONE; NEOMYCIN SULFATE; POLYMYXIN B SULFATE

SOLUTION/DROPS; OTIC

CORTISPORIN

+ MONARCH PHARMS

1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML **	N050479	001
--	---------	-----

NEOMYCIN AND POLYMYXIN B SULFATES AND HYDROCORTISONE

AMRING PHARMS

1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML	A065216	001	Oct 31, 2005
-------------------------------------	---------	-----	--------------

NEOMYCIN SULFATE-POLYMYXIN B SULFATE-HYDROCORTISONE

PHARMAFAIR

1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML	A062394	001	Sep 29, 1982
-------------------------------------	---------	-----	--------------

OTOCORT

WATSON LABS

1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML	A060730	002
-------------------------------------	---------	-----

SUSPENSION/DROPS; OPHTHALMIC

CORTISPORIN

MONARCH PHARMS

1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML	N050169	001
-------------------------------------	---------	-----

NEOMYCIN SULFATE-POLYMYXIN B SULFATE-HYDROCORTISONE

PHARMAFAIR

1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML	A062623	001	Sep 24, 1985
-------------------------------------	---------	-----	--------------

SUSPENSION/DROPS; OTIC

NEOMYCIN SULFATE, POLYMYXIN B SULFATE & HYDROCORTISONE

PHARMAFAIR

1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML	A062617	001	Sep 18, 1985
-------------------------------------	---------	-----	--------------

OTICAIR

PHARMAFAIR

1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML	A062399	001	Nov 18, 1982
-------------------------------------	---------	-----	--------------

OTOBIONE

SCHERING

1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML	A061816	001
-------------------------------------	---------	-----

OTOCORT

ACTAVIS LABS FL INC

1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML	A062521	001	Jul 11, 1985
-------------------------------------	---------	-----	--------------

PEDIOTIC

MONARCH PHARMS

1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML	A062822	001	Sep 29, 1987
-------------------------------------	---------	-----	--------------

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

HYDROCORTISONE; POLYMYXIN B SULFATE

SOLUTION/DROPS;OTIC

OTOBIOTIC

SCHERING 5MG/ML;EQ 10,000 UNITS BASE/ML A062302 001

PYOCIDIN

FOREST LABS 5MG/ML;EQ 10,000 UNITS BASE/ML A061606 001

HYDROCORTISONE; TETRACYCLINE HYDROCHLORIDE

OINTMENT;OPHTHALMIC

ACHROMYCIN

LEDERLE 1.5%;1% N050272 001

HYDROCORTISONE; UREA

CREAM;TOPICAL

ALPHADERM

BIOGLAN 1%;10% A086008 001

CALMURID HC

PHARMACIA AND UPJOHN 1%;10% A083947 001

HYDROFLUMETHIAZIDE

TABLET;ORAL

DIUCARDIN

WYETH AYERST 50MG A083383 001

HYDROFLUMETHIAZIDE

PAR PHARM 50MG A088850 001 May 31, 1985

WATSON LABS 50MG A088031 001 Apr 06, 1983

50MG A088528 001 Aug 15, 1984

SALURON

+ SHIRE LLC 50MG N011949 001

HYDROFLUMETHIAZIDE; RESERPINE

TABLET;ORAL

HYDROFLUMETHIAZIDE AND RESERPINE

USL PHARMA 50MG;0.125MG A088195 001 Oct 26, 1983

WATSON LABS 25MG;0.125MG A088127 001 Mar 22, 1983

50MG;0.125MG A088110 001 Mar 22, 1983

RESERPINE AND HYDROFLUMETHIAZIDE

IVAX PHARMS 50MG;0.125MG A088932 001 Jan 11, 1985

PAR PHARM 50MG;0.125MG A088907 001 Sep 20, 1985

SALUTENSIN

SHIRE 50MG;0.125MG N012359 003

SALUTENSIN-DEMI

SHIRE 25MG;0.125MG N012359 004

HYDROGEN PEROXIDE

SOLUTION;TOPICAL

ESKATA

+ ACLARIS 40% N209305 001 Dec 14, 2017

HYDROMORPHONE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE;ORAL

PALLADONE

PURDUE PHARMA LP 12MG N021044 001 Sep 24, 2004

16MG N021044 002 Sep 24, 2004

24MG N021044 003 Sep 24, 2004

32MG N021044 004 Sep 24, 2004

INJECTABLE;INJECTION

DILAUDID-HP

+ FRESENIUS KABI USA 10MG/ML N019034 001 Jan 11, 1984

250MG/VIAL N019034 002 Aug 04, 1994

HYDROMORPHONE HYDROCHLORIDE

BARR 10MG/ML A076444 001 Apr 25, 2003

HOSPIRA 10MG/ML A074598 001 Jun 19, 1997

WATSON LABS 10MG/ML A074317 001 Aug 23, 1995

SOLUTION;ORAL

HYDROMORPHONE HYDROCHLORIDE

GENUS LIFESCIENCES 5MG/5ML A207108 001 Apr 22, 2020

TABLET;ORAL

HYDROMORPHONE HYDROCHLORIDE

GENUS LIFESCIENCES 2MG A077471 002 Dec 09, 2009

4MG A077471 003 Dec 09, 2009

8MG A077471 001 Dec 09, 2009

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

HYDROMORPHONE HYDROCHLORIDE

TABLET; ORAL

HYDROMORPHONE HYDROCHLORIDE

HIKMA	4MG	A074597 003	May 29, 2009
	8MG	A074597 001	Jul 29, 1998
NESHER PHARMS	2MG	A077311 001	Nov 09, 2005
	4MG	A077311 002	Nov 09, 2005
	8MG	A077311 003	Nov 09, 2005
NOSTRUM LABS INC	8MG	A076723 001	Oct 18, 2005

TABLET, EXTENDED RELEASE; ORAL

EXALGO

+ SPECGX LLC	8MG **	N021217 001	Mar 01, 2010
+	12MG **	N021217 002	Mar 01, 2010
+	16MG **	N021217 003	Mar 01, 2010
+	32MG **	N021217 004	Aug 24, 2012

HYDROMORPHONE HYDROCHLORIDE

ACTAVIS LABS FL INC	8MG	A202144 001	May 12, 2014
	12MG	A202144 002	May 12, 2014
	16MG	A202144 003	May 12, 2014
	32MG	A202144 004	Jun 30, 2016

HYDROXOCOBALAMIN

INJECTABLE; INJECTION

ALPHAREDISOL

MERCK	1MG/ML	A080778 001	
-------	--------	-------------	--

CYANOKIT

BTG INTL	2.5GM/VIAL (5GM/KIT)	N022041 002	Dec 15, 2006
----------	----------------------	-------------	--------------

HYDROXOCOBALAMIN

ABRAXIS PHARM	1MG/ML	A084921 001	
WATSON LABS	1MG/ML	A085528 001	

HYDROXOMIN

BEL MAR	1MG/ML	A084629 001	
---------	--------	-------------	--

HYDROXYAMPHETAMINE HYDROBROMIDE

SOLUTION/DROPS; OPHTHALMIC

PAREDRIINE

PHARMICS	1%	N000004 004	
----------	----	-------------	--

HYDROXYAMPHETAMINE HYDROBROMIDE; TROPICAMIDE

SOLUTION/DROPS; OPHTHALMIC

PAREMYD

+ EPIC PHARMA LLC	1%; 0.25%	N019261 001	Jan 30, 1992
-------------------	-----------	-------------	--------------

HYDROXYCHLOROQUINE SULFATE

TABLET; ORAL

HYDROXYCHLOROQUINE SULFATE

HIKMA PHARMS	200MG	A040760 001	Aug 15, 2007
WATSON LABS	200MG	A040133 001	Nov 30, 1995

HYDROXYPROGESTERONE CAPROATE

INJECTABLE; INJECTION

HYDROXYPROGESTERONE CAPROATE

ALLERGAN	125MG/ML	N017439 001	
	250MG/ML	N017439 002	
EPIC PHARMA LLC	125MG/ML	N018004 001	

SOLUTION; INTRAMUSCULAR

DELALUTIN

+ BRISTOL MYERS SQUIBB	125MG/ML (125MG/ML) **	N010347 004	
+	125MG/ML (125MG/ML) **	N016911 001	
+	250MG/ML (250MG/ML) **	N010347 002	
+	250MG/ML (250MG/ML) **	N016911 002	

HYDROXYPROGESTERONE CAPROATE

ASPEN GLOBAL INC	1250MG/5ML (250MG/ML)	A200271 001	Aug 24, 2015
EUGIA PHARMA	1250MG/5ML (250MG/ML)	A211142 001	May 09, 2019

HYDROXYPROPYL CELLULOSE

INSERT; OPHTHALMIC

LACRISERT

+ BAUSCH AND LOMB INC	5MG	N018771 001	
-----------------------	-----	-------------	--

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

HYDROXYSTILBAMIDINE ISETHIONATE

INJECTABLE; INJECTION

HYDROXYSTILBAMIDINE ISETHIONATE

SANOFI AVENTIS US 225MG/AMP N009166 001

HYDROXYUREA

CAPSULE; ORAL

DROXIA

+	CHEPLAPHARM	200MG	N016295 002	Feb 25, 1998
+		300MG	N016295 003	Feb 25, 1998
+		400MG	N016295 004	Feb 25, 1998

HYDROXYUREA

	BARR	250MG	A075143 002	Sep 21, 2000
	BARR LABS INC	250MG	A075020 002	Jun 26, 2000
		500MG	A075020 001	Jul 30, 1998
	ROXANE	500MG	A074476 001	Aug 18, 1995

TABLET; ORAL

HYDROXYUREA

BARR 1GM A075734 001 Aug 29, 2000

HYDROXYZINE HYDROCHLORIDE

INJECTABLE; INJECTION

HYDROXYZINE

BAXTER HLTHCARE 50MG/ML A085551 002

HYDROXYZINE HYDROCHLORIDE

	ALTANA	25MG/ML	A087273 001	Apr 20, 1982
		50MG/ML	A087273 002	Apr 20, 1982
	BAXTER HLTHCARE	25MG/ML	A085551 001	
	DR REDDYS	50MG/ML	A085779 001	
	FRESENIUS KABI USA	25MG/ML	A087329 001	
		25MG/ML	A088184 001	Mar 31, 1983
		50MG/ML	A087329 002	
		50MG/ML	A088185 001	Mar 31, 1983
	HOSPIRA	25MG/ML	A087416 001	
		50MG/ML	A086821 001	
		50MG/ML	A087546 001	
	PHARMAFAIR	25MG/ML	A088862 001	Feb 14, 1986
		25MG/ML	A089106 001	Feb 14, 1986
		50MG/ML	A088881 001	Feb 14, 1986
		50MG/ML	A089107 001	Feb 14, 1986
	SMITH AND NEPHEW	25MG/ML	A087592 001	
	SOLOPAK	25MG/ML	A086822 001	
		25MG/ML	A087591 001	
		50MG/ML	A087310 001	
		50MG/ML	A087593 001	
		50MG/ML	A087595 001	
		50MG/ML	A087596 001	
	WATSON LABS	25MG/ML	A085778 001	
		25MG/ML	A087274 001	
		50MG/ML	A087274 002	
	WYETH AYERST	25MG/ML	A086258 001	
		50MG/ML	A086258 002	

ORGATRAK

	ORGANON USA INC	25MG/ML	A087014 001	
		50MG/ML	A087014 002	

VISTARIL

+	PFIZER	25MG/ML **	N011111 001	
+		50MG/ML **	N011111 002	

SYRUP; ORAL

ATARAX

ROERIG 10MG/5ML ** N010485 001

HYDROXYZINE HYDROCHLORIDE

	ALPHARMA US PHARMS	10MG/5ML	A088785 001	Feb 03, 1988
	ANIMA	10MG/5ML	A086880 001	
	HIKMA	10MG/5ML	A040010 001	Oct 28, 1994
	KV PHARM	10MG/5ML	A087730 001	Jul 01, 1982

TABLET; ORAL

ATARAX

+	PFIZER	10MG **	N010392 001	
+		25MG **	N010392 004	

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

HYDROXYZINE HYDROCHLORIDE

TABLET; ORAL

ATARAX

+

50MG **

N010392 006

+

100MG **

N010392 005

HYDROXYZINE HYDROCHLORIDE

ABLE

10MG

A040559 001 Jul 22, 2004

25MG

A040562 001 Jul 22, 2004

50MG

A040563 001 Jul 22, 2004

ACTAVIS ELIZABETH

10MG

A089071 001 Jul 22, 1986

25MG

A089072 001 Jul 22, 1986

50MG

A089073 001 Jul 22, 1986

HALSEY

10MG

A089366 001 May 02, 1988

25MG

A089117 001 May 02, 1988

50MG

A089396 001 May 02, 1988

INVAGEN PHARMS

10MG

A040812 001 Mar 12, 2008

25MG

A040812 002 Mar 12, 2008

50MG

A040812 003 Mar 12, 2008

IVAX PHARMS

10MG

A087216 001

25MG

A087410 001

50MG

A087411 001

KV PHARM

10MG

A087819 001 Jun 23, 1982

25MG

A087820 001 Jun 23, 1982

50MG

A087821 001 Jun 23, 1982

MUTUAL PHARM

100MG

A087822 001 Jun 23, 1982

10MG

A088409 001 Nov 15, 1983

25MG

A087857 001 Apr 18, 1983

50MG

A087860 001 Apr 18, 1983

PLIVA

100MG

A081054 001 Sep 25, 1995

PUREPAC PHARM

10MG

A088120 001 Sep 25, 1984

25MG

A088121 001 Sep 25, 1984

50MG

A088122 001 Sep 25, 1984

QUANTUM PHARMICS

10MG

A088540 001 Oct 22, 1985

25MG

A088551 001 Oct 22, 1985

50MG

A088529 001 Oct 22, 1985

RISING

10MG

A091176 001 Jun 07, 2010

25MG

A091176 002 Jun 07, 2010

50MG

A091176 003 Jun 07, 2010

SANDOZ

10MG

A087246 002

25MG

A085247 001

50MG

A087245 001

STRIDES PHARMA

10MG

A087602 001 Jan 22, 1982

25MG

A087603 001 Jan 22, 1982

50MG

A087604 001 Jan 22, 1982

SUN PHARM INDS INC

10MG

A040899 001 Jun 10, 2008

25MG

A040899 002 Jun 10, 2008

50MG

A040899 003 Jun 10, 2008

SUN PHARM INDUSTRIES

10MG

A089381 001 May 19, 1986

25MG

A089382 001 May 19, 1986

50MG

A089383 001 May 19, 1986

100MG

A087862 001 Apr 18, 1983

SUPERPHARM

10MG

A088794 001 Dec 05, 1984

25MG

A088795 001 Dec 05, 1984

50MG

A088796 001 Dec 05, 1984

USL PHARMA

10MG

A089121 001 Mar 20, 1986

25MG

A089122 001 Mar 20, 1986

50MG

A089123 001 Mar 20, 1986

WATSON LABS

10MG

A081149 001 Mar 18, 1994

10MG

A086827 001

10MG

A088348 001 Sep 15, 1983

25MG

A081150 001 Mar 18, 1994

25MG

A086829 001

25MG

A088349 001 Sep 15, 1983

50MG

A081151 001 Mar 18, 1994

50MG

A086836 001

50MG

A088350 001 Sep 15, 1983

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

HYDROXYZINE PAMOATE

CAPSULE;ORAL

HY-PAM "25"

TEVA EQ 25MG HYDROCHLORIDE A088713 001 Mar 04, 1985

HYDROXYZINE PAMOATE

BEXIMCO PHARMS USA EQ 25MG HYDROCHLORIDE A081127 001 Jun 28, 1991

DURAMED PHARMS BARR EQ 25MG HYDROCHLORIDE A088593 001 Feb 29, 1984

EQ 50MG HYDROCHLORIDE A088594 001 Feb 29, 1984

EQ 100MG HYDROCHLORIDE A088595 001 Feb 29, 1984

IVAX SUB TEVA PHARMS EQ 25MG HYDROCHLORIDE A087761 001 Mar 05, 1982

EQ 50MG HYDROCHLORIDE A087760 001 Mar 05, 1982

PAR PHARM EQ 25MG HYDROCHLORIDE A087656 001 Jun 11, 1982

EQ 25MG HYDROCHLORIDE A089145 001 Mar 17, 1986

EQ 50MG HYDROCHLORIDE A087657 001 Jun 11, 1982

EQ 50MG HYDROCHLORIDE A089146 001 Mar 17, 1986

EQ 100MG HYDROCHLORIDE A087658 001 Jun 11, 1982

SANDOZ EQ 50MG HYDROCHLORIDE A081128 001 Jun 28, 1991

EQ 100MG HYDROCHLORIDE A081129 001 Jun 28, 1991

SUPERPHARM EQ 25MG HYDROCHLORIDE A089031 001 Jan 02, 1987

EQ 50MG HYDROCHLORIDE A089032 001 Jan 02, 1987

EQ 100MG HYDROCHLORIDE A089033 001 Jan 02, 1987

VANGARD EQ 25MG HYDROCHLORIDE A088392 001 Sep 19, 1983

EQ 50MG HYDROCHLORIDE A088393 001 Sep 19, 1983

WATSON LABS EQ 25MG HYDROCHLORIDE A081165 001 Jul 31, 1991

EQ 25MG HYDROCHLORIDE A086698 001

EQ 25MG HYDROCHLORIDE A086840 001 Jul 01, 1982

EQ 50MG HYDROCHLORIDE A086695 001

EQ 50MG HYDROCHLORIDE A086705 001 Jul 01, 1982

EQ 50MG HYDROCHLORIDE A087767 001 Aug 16, 1982

EQ 100MG HYDROCHLORIDE A086697 001

EQ 100MG HYDROCHLORIDE A086728 001 Oct 05, 1982

EQ 100MG HYDROCHLORIDE A087790 001 Aug 16, 1982

VISTARIL

+ PFIZER

EQ 100MG HYDROCHLORIDE ** N011459 006

SUSPENSION;ORAL

VISTARIL

PFIZER

EQ 25MG HYDROCHLORIDE/5ML ** N011795 001

IBANDRONATE SODIUM

INJECTABLE;INTRAVENOUS

BONIVA

+ ROCHE

EQ 3MG BASE/3ML ** N021858 001 Jan 06, 2006

IBANDRONATE SODIUM

AVET LIFESCIENCES EQ 3MG BASE/3ML A203987 001 Sep 02, 2014

NANG KUANG PHARM CO EQ 3MG BASE/3ML A204329 001 Jun 16, 2021

SUN PHARM EQ 3MG BASE/3ML A090853 001 Feb 14, 2014

TABLET;ORAL

BONIVA

+ HOFFMANN LA ROCHE

EQ 2.5MG BASE ** N021455 001 May 16, 2003

+

EQ 150MG BASE ** N021455 002 Mar 24, 2005

IBANDRONATE SODIUM

NORVIUM BIOSCIENCE EQ 150MG BASE A078995 001 Mar 19, 2012

SUN PHARM INDUSTRIES EQ 150MG BASE A078996 001 Aug 15, 2012

WATSON LABS TEVA EQ 150MG BASE A079003 001 Mar 20, 2012

IBRUTINIB

CAPSULE;ORAL

IBRUTINIB

ZYDUS LIFESCIENCES 70MG A211344 001 Mar 31, 2021

140MG A211344 002 Mar 31, 2021

TABLET;ORAL

IMBRUVICA

+ PHARMACYCLICS LLC

560MG N210563 004 Feb 16, 2018

IBUPROFEN

CAPSULE;ORAL

IBUPROFEN

CONTRACT PHARMACAL 200MG A074782 001 Jul 06, 1998

MIDOL

BAYER 200MG ** A070626 001 Sep 02, 1987

200MG ** A071002 001 Sep 02, 1987

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

IBUPROFEN

SOLUTION; INTRAVENOUS

CALDOLOR

CUMBERLAND PHARMS 400MG/4ML (100MG/ML) N022348 001 Jun 11, 2009

SUSPENSION; ORAL

CHILDREN'S ADVIL

HALEON US HOLDINGS 100MG/5ML N019833 002 Sep 19, 1989

CHILDREN'S ELIXSURE

MOBERG PHARMA NORTH 100MG/5ML N021604 001 Jan 07, 2004

IBU

ABBOTT 100MG/5ML N019784 001 Dec 18, 1989

IBUPROFEN

ARISE 100MG/5ML A200457 001 Aug 18, 2011

PAI HOLDINGS PHARM 100MG/5ML A205647 001 Nov 03, 2016

STRIDES PHARMA 100MG/5ML A211666 001 Feb 22, 2021

MOTRIN

+ MCNEIL CONSUMER 100MG/5ML ** N019842 001 Sep 19, 1989

SUSPENSION/DROPS; ORAL

IBUPROFEN

AUROBINDO PHARMA LTD 40MG/ML A213506 001 Apr 08, 2024

STRIDES PHARMA 50MG/1.25ML A214071 001 Jun 09, 2022

MOTRIN

MCNEIL 40MG/ML N020476 001 May 25, 1995

PEDIATRIC ADVIL

+ HALEON US HOLDINGS 100MG/2.5ML N020812 001 Jan 30, 1998

TABLET; ORAL

ACHES-N-PAIN

LEDERLE 200MG A071065 001 May 28, 1987

CAP-PROFEN

PERRIGO 200MG A072097 001 Dec 08, 1987

IBU

BASF 400MG A070083 001 Feb 22, 1985

400MG N018197 001

600MG A070088 001 Feb 08, 1985

600MG A070099 001 Mar 29, 1985

800MG A070745 001 Jul 23, 1986

IBU-TAB

ALRA 400MG A071058 001 Aug 11, 1988

600MG A071059 001 Aug 11, 1988

800MG A071965 001 Aug 11, 1988

IBU-TAB 200

ALRA 200MG A071057 001 Aug 11, 1988

IBUPRIN

PLIVA 200MG A071773 001 Jul 16, 1987

IBUPROFEN

ABBOTT 600MG A070556 001 Jun 14, 1985

800MG A071264 001 Jul 25, 1986

ADAPTIS 200MG A071265 001 Oct 15, 1986

200MG A071265 002 Sep 10, 1987

ANI PHARMS 200MG A071144 001 Jan 20, 1987

200MG A072901 001 Dec 19, 1991

200MG A072903 001 Dec 19, 1991

CONTRACT PHARMACAL 200MG A071735 001 Sep 10, 1987

200MG A073691 001 Feb 25, 1994

200MG A074931 001 Jul 20, 1998

ENDO OPERATIONS 200MG A071639 001 Feb 02, 1988

200MG A072249 001 Jan 10, 1989

300MG A071230 001 Oct 22, 1986

400MG A071231 001 Oct 22, 1986

400MG A071644 001 Feb 01, 1988

600MG A071232 001 Oct 22, 1986

800MG A072004 001 Nov 18, 1987

HALSEY 200MG A071027 001 Sep 29, 1987

300MG A071028 001 Mar 23, 1987

400MG A071029 001 Mar 23, 1987

600MG A071030 001 Mar 23, 1987

800MG A072137 001 Feb 05, 1988

IVAX SUB TEVA PHARMS 200MG A071154 001 Oct 27, 1987

200MG A072040 001 Apr 29, 1988

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

IBUPROFEN

TABLET; ORAL

IBUPROFEN

	400MG	A071145	001	Sep 23, 1986
	600MG	A071146	001	Sep 23, 1986
	800MG	A071769	001	May 08, 1987
J AND J CONSUMER INC	400MG	A070081	001	Jun 16, 1986
L PERRIGO CO	400MG	A077114	001	Jul 18, 2005
	600MG	A077114	002	Jul 18, 2005
	800MG	A077114	003	Jul 18, 2005
LEDERLE	400MG	A070629	001	Sep 19, 1986
	600MG	A070630	001	Sep 19, 1986
LEINER	300MG	A071266	001	Oct 15, 1986
LNK	100MG	A076741	001	Jun 17, 2004
MCNEIL	600MG	A070476	001	Jun 16, 1986
MERRO PHARM	200MG	A070985	001	Oct 02, 1987
NORTHSTAR HLTHCARE	400MG	A078132	001	Sep 10, 2007
	600MG	A078132	002	Sep 10, 2007
	800MG	A078132	003	Sep 10, 2007
NORVIUM BIOSCIENCE	200MG	A071870	001	May 05, 1988
	400MG	A070045	001	Sep 24, 1985
	600MG	A070057	001	Sep 24, 1985
	800MG	A071999	001	Dec 03, 1987
OHM	200MG	A071163	001	Jul 15, 1986
OHM LABS	400MG	A070818	001	Dec 26, 1985
P AND L DEV LLC	200MG	A070733	001	Sep 19, 1986
PAR PHARM	300MG	A070328	001	Aug 06, 1985
PERRIGO	200MG	A072098	001	Dec 08, 1987
	200MG	A075995	001	Mar 14, 2002
PLIVA	400MG	A071666	001	Jun 18, 1987
	600MG	A071667	001	Jun 18, 1987
	800MG	A071668	001	Jun 18, 1987
PUREPAC PHARM	200MG	A071122	001	Oct 03, 1986
	200MG	A071664	001	Feb 03, 1987
	300MG	A071123	001	Sep 19, 1986
	400MG	A071124	001	Sep 19, 1986
	600MG	A071125	001	Sep 19, 1986
	800MG	A071964	001	Feb 01, 1988
RISING	300MG	A070736	002	Jun 12, 1986
	400MG	A070736	003	Jun 12, 1986
	600MG	A070736	001	Jun 12, 1986
	800MG	A071938	001	Jan 14, 1988
SANDOZ	200MG	A071807	001	Feb 25, 1988
	200MG	A074525	001	Dec 15, 1995
	200MG	A074533	001	Dec 15, 1995
	400MG	A072064	001	Jan 14, 1988
	600MG	A072065	001	Jan 14, 1988
	800MG	A072169	001	Dec 11, 1987
STRIDES PHARMA	200MG	A071575	001	May 08, 1987
	400MG	A070329	001	Aug 06, 1985
	600MG	A070330	001	Aug 06, 1985
	800MG	A070986	001	Jul 25, 1986
SUN PHARM INDUSTRIES	200MG	A070493	001	Dec 24, 1985
	200MG	A070908	001	Sep 26, 1986
	200MG	A071462	001	Oct 02, 1986
	400MG	A070079	001	Jul 24, 1985
	600MG	A070080	001	Jul 24, 1985
	800MG	A071448	001	Feb 18, 1987
SUNSHINE	400MG	A204062	001	Sep 10, 2018
	600MG	A204062	002	Sep 10, 2018
	800MG	A204062	003	Sep 10, 2018
SUPERPHARM	600MG	A070709	001	Apr 25, 1986
TEVA	200MG	A073141	001	May 29, 1992
	400MG	A073343	001	Jun 30, 1992
	600MG	A073344	001	Jun 30, 1992
	800MG	A073345	001	Jun 30, 1992
ULTRATAB LABS INC	200MG	A209076	001	Jan 06, 2020
WATSON LABS	200MG	A070435	001	Mar 05, 1986
	200MG	A071765	001	Sep 04, 1987

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

IBUPROFEN

TABLET;ORAL

IBUPROFEN

	200MG		A071905 001	Mar 08, 1988
	300MG		A071338 001	Dec 01, 1986
	400MG		A070038 001	Sep 06, 1985
	400MG		A070436 001	Aug 21, 1985
	600MG		A070041 001	Sep 06, 1985
	600MG		A070437 001	Aug 21, 1985
	800MG		A071547 001	Jul 02, 1987
	800MG		A071911 001	Oct 13, 1987
	200MG	YICHANG HUMANWELL	A214003 001	Oct 19, 2020
IBUPROHM				
	200MG	OHM LABS	A071214 001	Dec 01, 1986
	400MG		A070469 001	Aug 29, 1985
JUNIOR STRENGTH IBUPROFEN				
	100MG	L PERRIGO CO	A075367 001	Apr 22, 1999
MEDIPREN				
	200MG	MCNEIL	A070475 001	Feb 06, 1986
	200MG		A071215 001	Jun 26, 1986
MIDOL				
	200MG	BAYER	A070591 001	Sep 02, 1987
	200MG		A071001 001	Sep 02, 1987
MOTRIN				
	300MG **	+ MCNEIL CONSUMER	N017463 003	
	400MG **	+	N017463 002	
	600MG **	+	N017463 004	
	800MG **	+	N017463 005	May 22, 1985
	100MG	MCNEIL PED	N020418 001	Nov 16, 1994
MOTRIN MIGRAINE PAIN				
	200MG	KENVUE BRANDS	N019012 004	Feb 25, 2000
NUPRIN				
	200MG	BRISTOL MYERS	A072035 001	Feb 16, 1988
	200MG		A072036 001	Feb 16, 1988
	200MG	KENVUE BRANDS	N019012 001	May 18, 1984
	200MG		N019012 002	Jul 29, 1987
RUFEN				
	600MG	BASF	N018197 002	Mar 05, 1984
TAB-PROFEN				
	200MG	PERRIGO	A072095 001	Dec 08, 1987
TABLET, CHEWABLE;ORAL				
CHILDREN'S MOTRIN				
	50MG	+ KENVUE BRANDS	N020601 001	Nov 15, 1996
IBUPROFEN				
	50MG	PERRIGO	A076359 001	Jan 16, 2004
JUNIOR STRENGTH MOTRIN				
	100MG	+ KENVUE BRANDS	N020601 003	Nov 15, 1996
MOTRIN				
	50MG	MCNEIL PED	N020135 001	Nov 16, 1994
	100MG		N020135 002	Nov 16, 1994

IBUPROFEN SODIUM

TABLET;ORAL

IBUPROFEN SODIUM

PERRIGO R AND D EQ 200MG BASE

A206581 001 Aug 03, 2015

IBUPROFEN; OXYCODONE HYDROCHLORIDE

TABLET;ORAL

COMBUNOX

+ FOREST LABS 400MG;5MG **

N021378 001 Nov 26, 2004

OXYCODONE HYDROCHLORIDE AND IBUPROFEN

ACTAVIS ELIZABETH 400MG;5MG

A078769 001 Jan 04, 2008

BARR LABS INC 400MG;5MG

A078316 001 Nov 29, 2007

WATSON LABS 400MG;5MG

A078394 001 Nov 26, 2007

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

IBUPROFEN; PSEUDOEPHEDRINE HYDROCHLORIDE

SUSPENSION; ORAL

IBUPROFEN AND PSEUDOEPHEDRINE HYDROCHLORIDE

PERRIGO

100MG/5ML; 15MG/5ML

A076478 001 Nov 05, 2003

TABLET; ORAL

IBUPROFEN AND PSEUDOEPHEDRINE HYDROCHLORIDE

AUROBINDO PHARMA LTD 200MG; 30MG

A213565 001 Mar 10, 2023

CONTRACT PHARMACAL 200MG; 30MG

A075588 001 Apr 08, 2002

IBUTILIDE FUMARATE

INJECTABLE; INJECTION

IBUTILIDE FUMARATE

LUITPOLD

0.1MG/ML

A090240 001 Jan 11, 2010

MYLAN INSTITUTIONAL

0.1MG/ML

A090924 001 Jan 11, 2010

ICATIBANT ACETATE

INJECTABLE; SUBCUTANEOUS

ICATIBANT ACETATE

DR REDDYS

EQ 30MG BASE/3ML (EQ 10MG BASE/ML)

A213054 001 Oct 05, 2020

GLENMARK PHARMS LTD

EQ 30MG BASE/3ML (EQ 10MG BASE/ML)

A213222 001 May 21, 2021

IDARUBICIN HYDROCHLORIDE

INJECTABLE; INJECTION

IDAMYCIN

PFIZER

5MG/VIAL

N050661 002 Sep 27, 1990

10MG/VIAL

N050661 001 Sep 27, 1990

20MG/VIAL

N050661 003 Apr 25, 1995

IDARUBICIN HYDROCHLORIDE

FRESENIUS KABI USA 1MG/ML

A065440 001 Aug 04, 2009

RISING 1MG/ML

A200144 001 Oct 11, 2012

SANDOZ 1MG/ML

A091293 001 Mar 29, 2011

TEVA PARENTERAL 5MG/VIAL

A065037 003 May 01, 2002

10MG/VIAL

A065037 002 May 01, 2002

20MG/VIAL

A065037 001 May 01, 2002

IDOXURIDINE

OINTMENT; OPHTHALMIC

STOXIL

GLAXOSMITHKLINE

0.5%

N015868 001

SOLUTION/DROPS; OPHTHALMIC

DENDRID

+ ALCON

0.1%

N014169 001

HERPLEX

ALLERGAN

0.1%

N013935 002

STOXIL

GLAXOSMITHKLINE

0.1%

N013934 001

IFOSFAMIDE

INJECTABLE; INJECTION

IFOSFAMIDE

EXTROVIS

1GM/20ML (50MG/ML)

A201689 001 Nov 26, 2012

3GM/60ML (50MG/ML)

A201689 002 Nov 26, 2012

FRESENIUS KABI USA

1GM/20ML (50MG/ML)

A090181 001 Sep 22, 2009

3GM/60ML (50MG/ML)

A090181 002 Sep 22, 2009

IFOSFAMIDE; MESNA

INJECTABLE; INJECTION

IFEX/MESNEX KIT

BAXTER HLTHCARE

1GM/VIAL; 100MG/ML

N019763 003 Oct 10, 1992

3GM/VIAL; 100MG/ML

N019763 004 Oct 10, 1992

INJECTABLE; INTRAVENOUS

IFOSFAMIDE/MESNA KIT

TEVA PHARMS USA

1GM/20ML; 1GM/10ML (50MG/ML; 100MG/ML)

A075874 001 Feb 26, 2002

3GM/60ML; 1GM/10ML (50MG/ML; 100MG/ML)

A075874 002 Feb 26, 2002

ILOPERIDONE

TABLET; ORAL

ILOPERIDONE

INVENTIA

1MG

A207231 001 Nov 28, 2016

2MG

A207231 002 Nov 28, 2016

4MG

A207231 003 Nov 28, 2016

6MG

A207231 004 Nov 28, 2016

8MG

A207231 005 Nov 28, 2016

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ILOPERIDONE

TABLET; ORAL

ILOPERIDONE

	10MG	A207231 006	Nov 28, 2016
	12MG	A207231 007	Nov 28, 2016
LUPIN LTD	1MG	A206890 001	May 05, 2022
	2MG	A206890 002	May 05, 2022
	4MG	A206890 003	May 05, 2022
	6MG	A206890 004	May 05, 2022
	8MG	A206890 005	May 05, 2022
	10MG	A206890 006	May 05, 2022
	12MG	A206890 007	May 05, 2022
TARO	1MG	A207098 001	Jul 22, 2019
	2MG	A207098 002	Jul 22, 2019
	4MG	A207098 003	Jul 22, 2019
	6MG	A207098 004	Jul 22, 2019
	8MG	A207098 005	Jul 22, 2019
	10MG	A207098 006	Jul 22, 2019
	12MG	A207098 007	Jul 22, 2019

ILOPROST

SOLUTION; INHALATION

VENTAVIS

+	ACTELION	10MCG/ML (10MCG/ML)	N021779 002	Dec 08, 2005
		20MCG/2ML (10MCG/ML)	N021779 001	Dec 29, 2004
+		20MCG/ML (20MCG/ML)	N021779 003	Aug 07, 2009

IMATINIB MESYLATE

CAPSULE; ORAL

GLEEVEC

+	NOVARTIS	EQ 50MG BASE **	N021335 001	May 10, 2001
+		EQ 100MG BASE **	N021335 002	May 10, 2001

TABLET; ORAL

IMATINIB MESYLATE

AMNEAL PHARMS	EQ 100MG BASE	A207495 001	Feb 08, 2019
	EQ 400MG BASE	A207495 002	Feb 08, 2019
ESJAY PHARMA	EQ 100MG BASE	A205990 001	Feb 08, 2019
	EQ 400MG BASE	A205990 002	Feb 08, 2019
HIKMA	EQ 100MG BASE	A207586 001	Jul 13, 2018
	EQ 400MG BASE	A207586 002	Jul 13, 2018

IMIPRAMINE HYDROCHLORIDE

CONCENTRATE; ORAL

IMIPRAMINE HYDROCHLORIDE

NOVARTIS	25MG/ML	A086765 001	
----------	---------	-------------	--

INJECTABLE; INJECTION

TOFRANIL

NOVARTIS	12.5MG/ML	N011838 002	
----------	-----------	-------------	--

TABLET; ORAL

IMIPRAMINE HYDROCHLORIDE

CHARTWELL	10MG	A090441 002	Mar 11, 2010
	25MG	A090441 003	Mar 11, 2010
	50MG	A090441 001	Mar 11, 2010
LEDERLE	10MG	A086269 001	
	25MG	A086267 001	
	50MG	A086268 001	
PAR PHARM	10MG	A089422 001	Jul 14, 1987
	25MG	A089497 001	Jul 14, 1987
ROXANE	10MG	A083799 001	
	25MG	A083799 002	
	50MG	A083799 003	
SANDOZ	10MG	A085200 001	
	25MG	A084869 002	
	50MG	A085133 001	
+	SPECGX LLC	A087846 002	May 22, 1984
+		A087846 003	May 22, 1984
SUN PHARM INDUSTRIES	10MG	A081048 001	Jun 05, 1990
TEVA	10MG	A083729 001	
	25MG	A083729 004	
	50MG	A083729 003	
USL PHARMA	25MG	A087776 001	Feb 10, 1982

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

IMIPRAMINE HYDROCHLORIDE

TABLET; ORAL

IMIPRAMINE HYDROCHLORIDE

VANGARD	10MG	A088036 001	Nov 03, 1982
	25MG	A087619 001	Feb 09, 1982
	50MG	A087631 001	Jan 04, 1982
WATSON LABS	10MG	A085220 001	
	10MG	A085875 001	
	25MG	A084252 002	
	25MG	A085878 001	
	50MG	A085221 001	
	50MG	A085877 001	
WEST WARD	25MG	A088222 001	May 26, 1983
	50MG	A088223 001	May 26, 1983
JANIMINE			
ABBOTT	10MG	N017895 001	
	25MG	N017895 002	
	50MG	N017895 003	
PRAMINE			
ALRA	10MG	A083827 001	
	25MG	A083827 002	
	50MG	A083827 003	
PRESAMINE			
SANOFI AVENTIS US	10MG	N011836 006	
	25MG	N011836 003	
	50MG	N011836 007	
TOFRANIL			
+ SPECGX LLC	50MG	A087846 001	May 22, 1984

IMIPRAMINE PAMOATE

CAPSULE; ORAL

IMIPRAMINE PAMOATE

RISING	EQ 75MG HYDROCHLORIDE	A202338 001	Jun 28, 2013
	EQ 100MG HYDROCHLORIDE	A202338 002	Jun 28, 2013
	EQ 125MG HYDROCHLORIDE	A202338 003	Jun 28, 2013
	EQ 150MG HYDROCHLORIDE	A202338 004	Jun 28, 2013
TOFRANIL-PM			
+ SPECGX LLC	EQ 75MG HYDROCHLORIDE **	N017090 001	
	EQ 100MG HYDROCHLORIDE **	N017090 004	
	EQ 125MG HYDROCHLORIDE **	N017090 003	
	EQ 150MG HYDROCHLORIDE **	N017090 002	

IMIQUIMOD

CREAM; TOPICAL

ALDARA

+ BAUSCH	5% **	N020723 001	Feb 27, 1997
IMIQUIMOD			
COSETTE	5%	A200481 001	Apr 18, 2011
ENCUBE	5%	A091044 001	Feb 28, 2011
STRIDES PHARMA	5%	A202002 001	Jun 24, 2014

INAMRINONE LACTATE

INJECTABLE; INJECTION

AMRINONE LACTATE

BAXTER HLTHCARE CORP	EQ 5MG BASE/ML	A075542 001	May 10, 2000
HOSPIRA	EQ 5MG BASE/ML	A074616 001	Aug 03, 1998
INOCOR			
SANOFI AVENTIS US	EQ 5MG BASE/ML	N018700 001	Jul 31, 1984

INDACATEROL MALEATE

POWDER; INHALATION

ARCAPTA NEOHALER

+ NOVARTIS	EQ 75MCG BASE	N022383 001	Jul 01, 2011
------------	---------------	-------------	--------------

INDAPAMIDE

TABLET; ORAL

INDAPAMIDE

ANI PHARMS	1.25MG	A074498 002	Feb 12, 1998
	1.25MG	A075201 001	Dec 04, 1998
	2.5MG	A074498 001	Oct 31, 1996
	2.5MG	A075201 002	Dec 04, 1998
AUROBINDO PHARMA USA	1.25MG	A075105 001	Jul 23, 1998

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

INDAPAMIDETABLET; ORAL
INDAPAMIDE

	2.5MG	A075105 002	Jul 23, 1998
CHARTWELL RX	1.25MG	A074594 001	May 23, 1996
	2.5MG	A074594 002	May 23, 1996
TEVA	1.25MG	A074665 001	Apr 04, 1997
	2.5MG	A074665 002	Apr 04, 1997
WATSON LABS	1.25MG	A074585 001	Sep 26, 1996
	2.5MG	A074585 002	Sep 26, 1996
LOZOL			
+ SANOFI AVENTIS US	1.25MG **	N018538 002	Apr 29, 1993
+	2.5MG **	N018538 001	Jul 06, 1983

INDECAINIDE HYDROCHLORIDETABLET, EXTENDED RELEASE; ORAL
DECABID

LILLY	EQ 50MG BASE	N019693 001	Dec 29, 1989
	EQ 75MG BASE	N019693 002	Dec 29, 1989
	EQ 100MG BASE	N019693 003	Dec 29, 1989

INDINAVIR SULFATE

CAPSULE; ORAL

CRIXIVAN

MERCK SHARP DOHME	EQ 100MG BASE	N020685 006	Apr 19, 2000
+	EQ 200MG BASE	N020685 003	Mar 13, 1996
	EQ 333MG BASE	N020685 005	Dec 17, 1998
+	EQ 400MG BASE	N020685 001	Mar 13, 1996

INDIUM IN-111 CHLORIDE

INJECTABLE; INJECTION

INDICLOR

+ GE HEALTHCARE	2mCi/0.2ML	N019862 001	Dec 29, 1992
-----------------	------------	-------------	--------------

INDIUM IN 111 CHLORIDE

+ CURIUM	5mCi/0.5ML	N019841 001	Sep 27, 1994
----------	------------	-------------	--------------

INDOCYANINE GREEN

INJECTABLE; INJECTION

IC-GREEN

RENEW PHARMS	10MG/VIAL **	N011525 003	
+	25MG/VIAL	N011525 001	
	40MG/VIAL **	N011525 004	
	50MG/VIAL **	N011525 002	

INDOMETHACIN

CAPSULE; ORAL

INDO-LEMMON

TEVA	25MG	A070266 001	Nov 07, 1985
	50MG	A070267 001	Nov 07, 1985

INDOCIN

+ ZYLA LIFE SCIENCES	25MG **	N016059 001	
----------------------	---------	-------------	--

+	50MG **	N016059 002	
---	---------	-------------	--

INDOMETHACIN

ABLE	25MG	A076666 001	Dec 17, 2003
	50MG	A076666 002	Dec 17, 2003
ANI PHARMS	25MG	A071148 001	Mar 18, 1987
	50MG	A071149 001	Mar 18, 1987
CYCLE	25MG	A070353 001	Jun 18, 1985
	50MG	A070354 001	Jun 18, 1985
DURAMED PHARMS BARR	25MG	A070326 001	Oct 18, 1985
	50MG	A070327 001	Oct 18, 1985
HALSEY	25MG	A070782 001	Jun 03, 1987
	50MG	A070635 001	Jun 03, 1987
HERITAGE	25MG	N018851 001	May 18, 1984
	50MG	N018851 002	May 18, 1984
HERITAGE PHARMA	25MG	A070719 001	Feb 12, 1986
	50MG	A070756 001	Feb 12, 1986
IVAX SUB TEVA PHARMS	25MG	N018730 001	May 04, 1984
	50MG	N018730 002	May 04, 1984
JUBILANT GENERICS	25MG	A205215 001	Aug 25, 2017
	50MG	A205215 002	Aug 25, 2017
MUTUAL PHARM	25MG	A070067 001	Oct 03, 1986

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

INDOMETHACIN

CAPSULE; ORAL

INDOMETHACIN

	50MG	A070068 001 Oct 03, 1986
PARKE DAVIS	25MG	N018806 001 Nov 23, 1984
	50MG	N018806 002 Nov 23, 1984
PIONEER PHARMS	25MG	A070813 001 Aug 11, 1986
	50MG	A070592 001 Aug 11, 1986
RISING	25MG	N018858 001 Apr 20, 1984
	50MG	A070624 001 Sep 04, 1985
	50MG	N018858 002 Apr 20, 1984
SUN PHARM INDS INC	25MG	A091401 001 Mar 28, 2013
	50MG	A091401 002 Mar 28, 2013
SUN PHARM INDUSTRIES	25MG	A070900 002 Feb 09, 1987
	50MG	A070900 001 Feb 09, 1987
SUPERPHARM	25MG	A070487 001 Oct 10, 1986
	50MG	A070488 001 Oct 10, 1986
TEVA	25MG	A071342 001 Apr 18, 1988
	50MG	A071343 001 Apr 18, 1988
WATSON LABS	25MG	A070529 001 Oct 18, 1985
	25MG	A070784 001 Aug 20, 1986
	25MG	A072996 001 Jul 31, 1991
	25MG	N018690 001 Jul 31, 1984
	50MG	A070530 001 Oct 18, 1985
	50MG	A070785 001 Aug 20, 1986
	50MG	A071635 001 May 18, 1987
	50MG	A072997 001 Jul 31, 1991
	50MG	N018690 002 Jul 31, 1984

TIVORBEX

+ GENUS

20MG

N204768 001 Feb 24, 2014

+

40MG

N204768 002 Feb 24, 2014

CAPSULE, EXTENDED RELEASE; ORAL

INDOCIN SR

+ ZYLA

75MG **

N018185 001 Feb 23, 1982

INDOMETHACIN

ABLE

75MG

A076114 001 Feb 06, 2002

AUROBINDO PHARMA

75MG

A204243 001 Dec 27, 2016

INWOOD LABS

75MG

A072410 001 Mar 15, 1989

JUBILANT GENERICS

75MG

A202706 001 Oct 05, 2015

MYLAN

75MG

A202139 001 Mar 20, 2014

SANDOZ

75MG

A074464 001 May 28, 1998

WATSON LABS INC

75MG

A202572 001 Dec 09, 2013

SUPPOSITORY; RECTAL

INDOCIN

+ ZYLA LIFE SCIENCES

50MG **

N017814 001 Aug 13, 1984

SUSPENSION; ORAL

INDOMETHACIN

HIKMA

25MG/5ML

A071412 001 Mar 18, 1987

INDOMETHACIN SODIUM

INJECTABLE; INJECTION

INDOCIN

+ RECORDATI RARE

EQ 1MG BASE/VIAL **

N018878 001 Jan 30, 1985

INDOMETHACIN SODIUM

HIKMA

EQ 1MG BASE/VIAL

A078713 001 Jul 16, 2008

HOSPIRA

EQ 1MG BASE/VIAL

A204118 001 Apr 19, 2016

NAVINTA LLC

EQ 1MG BASE/VIAL

A206561 001 Jul 19, 2017

INFIGRATINIB PHOSPHATE

CAPSULE; ORAL

TRUSELTIQ

+ HELSINN HLTHCARE

25MG

N214622 001 May 28, 2021

+

100MG

N214622 002 May 28, 2021

INGENOL MEBUTATE

GEL; TOPICAL

INGENOL MEBUTATE

PADAGIS ISRAEL

0.015%

A209018 001 Jan 07, 2019

0.05%

A209019 001 Jan 09, 2019

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

INGENOL MEBUTATE

GEL; TOPICAL

PICATO

+ LEO LABS

0.015%

N202833 001 Jan 23, 2012

+

0.05%

N202833 002 Jan 23, 2012

INOTERSEN SODIUM

SOLUTION; SUBCUTANEOUS

TEGSEDI

+ AKCEA THERAPS

EQ 284MG BASE/1.5ML (EQ 189.3MG
BASE/ML)

N211172 001 Oct 05, 2018

INULIN

INJECTABLE; INJECTION

INULIN AND SODIUM CHLORIDE

ISO TEX

100MG/ML

N002282 001

INVERT SUGAR

INJECTABLE; INJECTION

TRAVERT 10% IN PLASTIC CONTAINER

BAXTER HLTHCARE

10GM/100ML

N016717 001

IOBENGUANE I-131

SOLUTION; INTRAVENOUS

AZEDRA

+ PROGENICS PHARMS INC 15mCi/ML

N209607 001 Jul 30, 2018

IOBENGUANE SULFATE I-131

INJECTABLE; INJECTION

IOBENGUANE SULFATE I 131

PHARMALUCENCE

2.3mCi/ML

N020084 001 Mar 25, 1994

IOCETAMIC ACID

TABLET; ORAL

CHOLEBRINE

MALLINCKRODT

750MG

N017129 001

IODAMIDE MEGLUMINE

INJECTABLE; INJECTION

RENOVUE-65

BRACCO

65%

N017902 001

RENOVUE-DIP

BRACCO

24%

N017903 001

IODIPAMIDE MEGLUMINE

INJECTABLE; INJECTION

CHOLOGRAFIN MEGLUMINE

BRACCO

10.3%

N009321 007

+

52%

N009321 003

IODIPAMIDE SODIUM

INJECTABLE; INJECTION

CHOLOGRAFIN SODIUM

BRACCO

20%

N009321 001

IODIXANOL

INJECTABLE; INJECTION

VISIPAQUE 270

GE HEALTHCARE

55%

N020808 001 Aug 29, 1997

IODOHIPPURATE SODIUM I-123

INJECTABLE; INJECTION

NEPHROFLOW

GE HEALTHCARE

1mCi/ML

N018289 001 Dec 28, 1984

IODOHIPPURATE SODIUM I-131

INJECTABLE; INJECTION

HIPURAN I 131

MALLINCKRODT

0.25mCi/ML

N016666 001

HIPPUTOPE

BRACCO

1-2mCi/VIAL

N015419 002

IODOHIPPURATE SODIUM I 131

PHARMALUCENCE

0.2mCi/ML

N017313 001

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

IODOXAMATE MEGLUMINE

INJECTABLE; INJECTION

CHOLOVUE

BRACCO	9.9%	N018077 001
	40.3%	N018076 001

IOFETAMINE HYDROCHLORIDE I-123

INJECTABLE; INJECTION

SPECTAMINE

IMP	1mCi/ML	N019432 001	Dec 24, 1987
-----	---------	-------------	--------------

IOHEXOL

FOR SOLUTION; ORAL

ORALTAG

INTERPHARMA PRAHA AS	9.7GM/BOT	N205383 001	Mar 26, 2015
----------------------	-----------	-------------	--------------

INJECTABLE; INJECTION

OMNIPAQUE 210

GE HEALTHCARE	45.3%	N018956 006	Jun 30, 1989
---------------	-------	-------------	--------------

SOLUTION; INJECTION, ORAL, RECTAL

OMNIPAQUE 240

GE HEALTHCARE	51.8%	N020608 001	Oct 24, 1995
---------------	-------	-------------	--------------

SOLUTION; URETHRAL

OMNIPAQUE 70

GE HEALTHCARE	15.1%	N018956 007	Jun 01, 1994
---------------	-------	-------------	--------------

IOPAMIDOL

INJECTABLE; INJECTION

IOPAMIDOL

BAXTER HLTHCARE	41%	A074629 001	Nov 06, 1996
	51%	A074629 004	Mar 31, 1998
	61%	A074629 002	Nov 06, 1996
	76%	A074629 003	Nov 06, 1996
HOSPIRA	61%	A074734 001	Dec 10, 1996
	76%	A074734 002	Dec 10, 1996

IOPAMIDOL-200

COOK IMAGING	41%	A074881 001	Jul 28, 2000
--------------	-----	-------------	--------------

HOSPIRA	41%	A074898 001	Dec 30, 1997
---------	-----	-------------	--------------

IOPAMIDOL-200 IN PLASTIC CONTAINER

HOSPIRA	41%	A074636 001	Dec 30, 1997
---------	-----	-------------	--------------

IOPAMIDOL-250

COOK IMAGING	51%	A074881 002	Jul 28, 2000
--------------	-----	-------------	--------------

FRESENIUS KABI USA	51%	A074679 001	Apr 02, 1997
--------------------	-----	-------------	--------------

HOSPIRA	51%	A074898 002	Dec 30, 1997
---------	-----	-------------	--------------

	51%	A075005 001	Feb 24, 1998
--	-----	-------------	--------------

IOPAMIDOL-250 IN PLASTIC CONTAINER

HOSPIRA	51%	A074636 002	Dec 30, 1997
---------	-----	-------------	--------------

IOPAMIDOL-300

ABBVIE	61%	A074638 001	Apr 30, 1997
--------	-----	-------------	--------------

COOK IMAGING	61%	A074881 003	Jul 28, 2000
--------------	-----	-------------	--------------

FRESENIUS KABI USA	61%	A074679 002	Apr 02, 1997
--------------------	-----	-------------	--------------

HOSPIRA	61%	A074898 003	Dec 30, 1997
---------	-----	-------------	--------------

	61%	A075005 002	Feb 24, 1998
--	-----	-------------	--------------

IOPAMIDOL-300 IN PLASTIC CONTAINER

HOSPIRA	61%	A074636 003	Dec 30, 1997
---------	-----	-------------	--------------

	61%	A074637 001	Apr 03, 1997
--	-----	-------------	--------------

IOPAMIDOL-370

COOK IMAGING	76%	A074881 004	Jul 28, 2000
--------------	-----	-------------	--------------

FRESENIUS KABI USA	76%	A074679 003	Apr 02, 1997
--------------------	-----	-------------	--------------

HOSPIRA	76%	A074898 004	Dec 30, 1997
---------	-----	-------------	--------------

	76%	A075005 003	Feb 24, 1998
--	-----	-------------	--------------

IOPAMIDOL-370 IN PLASTIC CONTAINER

HOSPIRA	76%	A074636 004	Dec 30, 1997
---------	-----	-------------	--------------

ISOVUE-128

BRACCO	26%	N018735 005	Oct 21, 1986
--------	-----	-------------	--------------

ISOVUE-200

BRACCO	41%	N020327 001	Oct 12, 1994
--------	-----	-------------	--------------

ISOVUE-250

+ BRACCO	51%	N020327 002	Oct 12, 1994
----------	-----	-------------	--------------

SCANLUX-300

SANOCHEMIA CORP USA	61%	A090394 001	Jun 18, 2010
---------------------	-----	-------------	--------------

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

IOPAMIDOL

INJECTABLE; INJECTION

SCANLUX-370

SANOCHEMIA CORP USA 76%

A090394 002 Jun 18, 2010

IOPANOIC ACID

TABLET; ORAL

TELEPAQUE

GE HEALTHCARE 500MG

N008032 001

IOPHENDYLATE

INJECTABLE; INJECTION

PANTOPAQUE

ALCON 100%

N005319 001

IOPROMIDE

INJECTABLE; INJECTION

ULTRAVIST (PHARMACY BULK)

+ BAYER HLTHCARE 49.9%

N021425 003 Mar 12, 2004

ULTRAVIST 150

+ BAYER HLTHCARE 31.2%

N020220 004 May 10, 1995

ULTRAVIST 240

+ BAYER HLTHCARE 49.9%

N020220 003 May 10, 1995

ULTRAVIST 300 IN PLASTIC CONTAINER

+ BAYER HLTHCARE 62.3%

N020220 005 Nov 18, 2008

IOTHALAMATE MEGLUMINE

INJECTABLE; INJECTION

CONRAY 30

+ LIEBEL-FLARSHEIM 30%

N016983 001

CONRAY 43

+ LIEBEL-FLARSHEIM 43%

N013295 002

IOTHALAMATE MEGLUMINE; IOTHALAMATE SODIUM

INJECTABLE; INJECTION

VASCORAY

MALLINCKRODT 52%;26%

N016783 001

IOTHALAMATE SODIUM

INJECTABLE; INJECTION

ANGIO-CONRAY

MALLINCKRODT 80%

N013319 001

CONRAY 325

MALLINCKRODT 54.3%

N017685 001

CONRAY 400

MALLINCKRODT 66.8%

N014295 001

IOTROLAN

INJECTABLE; INTRATHECAL

OSMOVIST 190

BAYER HLTHCARE 40.6%

N019580 001 Dec 07, 1989

OSMOVIST 240

BAYER HLTHCARE 51.3%

N019580 002 Dec 07, 1989

IOVERSOL

INJECTABLE; INJECTION

OPTIRAY 160

LIEBEL-FLARSHEIM 34%

N019710 003 Dec 30, 1988

OPTIRAY 240

+ LIEBEL-FLARSHEIM 51%

N019710 002 Dec 30, 1988

51%

N020923 001 May 28, 1998

IOXAGLATE MEGLUMINE; IOXAGLATE SODIUM

INJECTABLE; INJECTION

HEXABRIX

GUERBET 39.3%;19.6%

N018905 002 Jul 26, 1985

IOXILAN

INJECTABLE; INJECTION

OXILAN-300

GUERBET 62%

N020316 001 Dec 21, 1995

OXILAN-350

GUERBET 73%

N020316 002 Dec 21, 1995

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

IPODATE CALCIUM

GRANULE; ORAL

ORAGRAFIN CALCIUM

BRACCO

3GM/PACKET

N012968 001

IPODATE SODIUM

CAPSULE; ORAL

BILIVIST

BAYER HLTHCARE

500MG

A087768 001 Aug 11, 1982

ORAGRAFIN SODIUM

BRACCO

500MG

N012967 001

IPRATROPIUM BROMIDE

AEROSOL, METERED; INHALATION

ATROVENT

BOEHRINGER INGELHEIM 0.018MG/INH

N019085 001 Dec 29, 1986

SOLUTION; INHALATION

ATROVENT

+ BOEHRINGER INGELHEIM 0.02% **

N020228 001 Sep 29, 1993

IPRATROPIUM BROMIDE

ACTAVIS MID ATLANTIC 0.02%

A075111 001 Apr 22, 1999

APOTEX INC 0.02%

A075441 001 Mar 28, 2001

BAUSCH 0.02%

A075835 001 Oct 15, 2001

LANDELA PHARM 0.02%

A077072 001 Jul 19, 2005

NORVIUM BIOSCIENCE 0.02%

A074755 001 Jan 10, 1997

ROXANE 0.02%

A075867 001 Jul 22, 2002

TEVA PHARMS USA 0.02%

A075313 001 Feb 07, 2000

WATSON LABS 0.02%

A076291 001 May 09, 2005

ZENNOVA 0.02%

A075507 001 Jan 19, 2001

SPRAY, METERED; NASAL

ATROVENT

+ BOEHRINGER INGELHEIM 0.021MG/SPRAY **

N020393 001 Oct 20, 1995

+ 0.042MG/SPRAY **

N020394 001 Oct 20, 1995

IPRATROPIUM BROMIDE

AUROBINDO PHARMA USA 0.021MG/SPRAY

A075552 001 Mar 31, 2003

0.042MG/SPRAY

A075553 001 Mar 31, 2003

IRBESARTAN

TABLET; ORAL

AVAPRO

+ SANOFI AVENTIS US 75MG

N020757 001 Sep 30, 1997

IRBESARTAN

AJANTA PHARMA LTD 75MG

A203685 001 Dec 10, 2015

150MG

A203685 002 Dec 10, 2015

300MG

A203685 003 Dec 10, 2015

APOTEX INC 75MG

A200832 001 Oct 15, 2012

150MG

A200832 002 Oct 15, 2012

300MG

A200832 003 Oct 15, 2012

APPCO 75MG

A200461 001 Sep 27, 2012

150MG

A200461 002 Sep 27, 2012

300MG

A200461 003 Sep 27, 2012

CHARTWELL RX 75MG

A203161 001 Sep 27, 2012

150MG

A203161 002 Sep 27, 2012

300MG

A203161 003 Sep 27, 2012

HIKMA 75MG

A090201 001 Oct 15, 2012

150MG

A090201 002 Oct 15, 2012

300MG

A090201 003 Oct 15, 2012

IPCA LABS LTD 75MG

A211056 001 May 22, 2024

150MG

A211056 002 May 22, 2024

300MG

A211056 003 May 22, 2024

WATSON LABS INC 75MG

A090720 001 Oct 12, 2012

150MG

A090720 002 Oct 12, 2012

300MG

A090720 003 Oct 12, 2012

IRINOTECAN HYDROCHLORIDE

INJECTABLE; INJECTION

IRINOTECAN HYDROCHLORIDE

CHARTWELL RX 40MG/2ML (20MG/ML)

A078953 001 Apr 15, 2010

100MG/5ML (20MG/ML)

A078953 002 Apr 15, 2010

CIPLA LTD 40MG/2ML (20MG/ML)

A077219 001 Feb 20, 2008

100MG/5ML (20MG/ML)

A077219 002 Feb 20, 2008

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

IRINOTECAN HYDROCHLORIDE

INJECTABLE; INJECTION

IRINOTECAN HYDROCHLORIDE

EMCURE PHARMS LTD	40MG/2ML (20MG/ML)	A200771 001	Feb 14, 2012
	100MG/5ML (20MG/ML)	A200771 002	Feb 14, 2012
FRESENIUS KABI USA	40MG/2ML (20MG/ML)	A078188 001	Feb 27, 2008
	100MG/5ML (20MG/ML)	A078188 002	Feb 27, 2008
HISUN PHARM HANGZHOU	40MG/2ML (20MG/ML)	A090016 001	Jan 28, 2009
	100MG/5ML (20MG/ML)	A090016 002	Jan 28, 2009
PLIVA LACHEMA	40MG/2ML (20MG/ML)	A078122 001	Oct 31, 2008
	100MG/5ML (20MG/ML)	A078122 002	Oct 31, 2008
SANDOZ	40MG/2ML (20MG/ML)	A077994 001	Feb 27, 2008
	40MG/2ML (20MG/ML)	A090137 001	Nov 12, 2009
	100MG/5ML (20MG/ML)	A077994 002	Feb 27, 2008
	100MG/5ML (20MG/ML)	A090137 002	Nov 12, 2009
SUN PHARMA GLOBAL	40MG/2ML (20MG/ML)	A078805 001	Apr 21, 2008
	100MG/5ML (20MG/ML)	A078805 002	Apr 21, 2008
TEVA PHARMS USA	40MG/2ML (20MG/ML)	A090101 002	Feb 27, 2008
	100MG/5ML (20MG/ML)	A090101 003	Feb 27, 2008

ISOETHARINE HYDROCHLORIDE

SOLUTION; INHALATION

BETA-2

NEPHRON	1%	A086711 001	
BRONKOSOL			
SANOFI AVENTIS US	0.25%	N012339 009	
	1%	N012339 008	
ISOETHARINE HYDROCHLORIDE			
ALPHARMA US PHARMS	1%	A087101 001	
+ ASTRAZENECA	0.062%	A087937 001	Nov 15, 1982
	0.062%	A089614 001	Jun 13, 1991
+	0.125%	A087938 001	Nov 15, 1982
	0.125%	A089615 001	Jun 13, 1991
+	0.167%	A088470 001	Mar 14, 1984
	0.167%	A089616 001	Jun 13, 1991
+	0.2%	A088471 001	Mar 14, 1984
	0.2%	A089617 001	Jun 13, 1991
+	0.25%	A088472 001	Mar 14, 1984
	0.25%	A089618 001	Jun 13, 1991
BAXTER HLTHCARE	0.08%	A088144 001	Jul 29, 1983
	0.14%	A088145 001	Mar 26, 1984
	0.25%	A088146 001	Aug 01, 1983
DEY	0.08%	A088187 001	Dec 03, 1982
	0.1%	A087389 001	
	0.17%	A087390 001	
	0.25%	A088188 001	Dec 03, 1982
+	1%	A086763 001	
INTL MEDICATION	0.077%	A086651 001	
	0.08%	A086651 002	
	0.1%	A086651 003	
	0.143%	A086651 004	
	0.167%	A086651 005	
	0.2%	A086651 006	
	0.25%	A086651 007	
	1%	A086651 008	
PARKE DAVIS	0.5%	A085997 001	
	1%	A085889 001	
ROXANE	0.1%	A087396 001	
	0.125%	A087025 001	
	0.167%	A088226 001	Sep 16, 1983
	0.2%	A087324 001	
	0.25%	A088275 001	Jun 03, 1983
	1%	A086899 001	
ISOETHARINE HYDROCHLORIDE S/F			
DEY	0.08%	A089817 001	Nov 22, 1988
	0.1%	A089818 001	Nov 22, 1988
	0.17%	A089819 001	Nov 22, 1988
	0.25%	A089820 001	Nov 22, 1988
	1%	A089252 001	Sep 15, 1986

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ISOETHARINE MESYLATEAEROSOL, METERED; INHALATION
BRONKOMETER

SANOFI AVENTIS US 0.34MG/INH

N012339 007

ISOETHARINE MESYLATE

ALPHARMA US PHARMS 0.34MG/INH

A087858 001 Aug 21, 1984

ISOFLURANE

LIQUID; INHALATION

ISOFLURANE

HOSPIRA 99.9%

A074097 001 Jan 25, 1993

WATSON LABS INC 99.9%

A074393 001 May 12, 1995

ISOFLUROPHATE

OINTMENT; OPHTHALMIC

FLOROPRYL

MERCK 0.025%

N010656 001

ISONIAZID

INJECTABLE; INJECTION

NYDRAZID

SANDOZ 100MG/ML **

N008662 001

RIMIFON

ROCHE 25MG/ML

N008420 002

100MG/ML

N008420 003

SYRUP; ORAL

ISONIAZID

CHARTWELL RX 50MG/5ML

A081118 001 Jul 21, 1997

LANIAZID

LANNETT 50MG/5ML

A089243 001 Feb 03, 1986

RIMIFON

ROCHE 50MG/5ML

N008420 001

TABLET; ORAL

DOW-ISONIAZID

DOW PHARM 300MG

A080330 002

HYZYD

MEDPOINTE PHARM HLC 100MG

A080134 003

300MG

A080134 004

INH

NOVARTIS 300MG

A080935 001

ISONIAZID

DURAMED PHARMS BARR 100MG

A088231 001 Mar 17, 1983

300MG

A088119 001 Mar 17, 1983

+ EPIC PHARMA LLC 100MG **

N008678 002

+ 300MG **

N008678 003

HALSEY 50MG

A083632 001

HIKMA INTL PHARMS 100MG

A080212 001

300MG

A087425 001

IMPAX LABS 100MG

A080153 001

IVAX SUB TEVA PHARMS 100MG

A080270 001

300MG

A083610 001

LILLY 100MG

N008499 002

300MG

N008499 003

MK LABS 100MG

A080941 001

NEXGEN PHARMA INC 100MG

A084050 001

PANRAY 50MG

N008428 001

100MG

N008428 002

300MG

N008428 003

PERRIGO 100MG

A083060 001

PHARMAVITE 100MG

A085091 001

PHOENIX LABS NY 50MG

A080368 001

100MG

A080368 002

PUREPAC PHARM 50MG

A080132 003 Jul 14, 1982

100MG

A080132 004 Jul 14, 1982

SUN PHARM INDUSTRIES 100MG

A080136 001

300MG

A083633 001

WATSON LABS 50MG

A080522 001

100MG

A080401 001

100MG

A080523 001

100MG

A085790 001

300MG

A080521 001

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ISONIAZID

TABLET; ORAL

ISONIAZID

300MG	A083178	001
300MG	A085784	001
WHITEWORTH TOWN PLSN 100MG	A080120	002

LANIAZID

CHARTWELL MOLECULAR 50MG	A080140	001
100MG	A080140	002
300MG	A089776	001

Jun 13, 1988

NYDRAZID

BRISTOL MYERS SQUIBB 100MG	N008392	003
----------------------------	---------	-----

STANOZIDE

EVERYLIFE 100MG	A080126	001
300MG	A080126	002

ISONIAZID; PYRAZINAMIDE; RIFAMPIN

TABLET; ORAL

RIFATER

+ SANOFI AVENTIS US 50MG;300MG;120MG	N050705	001	May 31, 1994
--------------------------------------	---------	-----	--------------

ISONIAZID; RIFAMPIN

CAPSULE; ORAL

RIFAMATE

SANOFI AVENTIS US 150MG;300MG	A061884	001
-------------------------------	---------	-----

RIFAMPIN AND ISONIAZID

HIKMA INTL PHARMS 150MG;300MG	A065221	001	Jul 29, 2005
-------------------------------	---------	-----	--------------

ISOPROPAMIDE IODIDE

TABLET; ORAL

DARBID

GLAXOSMITHKLINE EQ 5MG BASE	N010744	001
-----------------------------	---------	-----

ISOPROTERENOL HYDROCHLORIDE

AEROSOL, METERED; INHALATION

ISOPROTERENOL HYDROCHLORIDE

3M 0.12MG/INH	N010375	004
ALPHARMA US PHARMS 0.12MG/INH	A085904	001

ISUPREL

SANOFI AVENTIS US 0.103MG/INH	N011178	001
-------------------------------	---------	-----

DISC; INHALATION

NORISODRINE AEROTROL

ABBOTT 0.25%	N016814	001
--------------	---------	-----

INJECTABLE; INJECTION

ISOPROTERENOL HYDROCHLORIDE

ABRAXIS PHARM 0.2MG/ML	A083431	001
------------------------	---------	-----

AM REGENT 0.2MG/ML	A214775	001	Oct 19, 2022
--------------------	---------	-----	--------------

BAXTER HLTHCARE 0.2MG/ML	A083486	001
--------------------------	---------	-----

CIPLA 0.2MG/ML	A210322	001	Jun 12, 2018
----------------	---------	-----	--------------

0.2MG/ML	A211738	001	Jun 28, 2019
----------	---------	-----	--------------

EUGIA PHARMA 0.2MG/ML	A211864	001	Feb 09, 2021
-----------------------	---------	-----	--------------

HOSPIRA 0.02MG/ML	A083283	001
-------------------	---------	-----

0.2MG/ML	A083346	001
----------	---------	-----

INTL MEDICATION 0.2MG/ML	A083724	001
--------------------------	---------	-----

NORVIUM BIOSCIENCE 0.2MG/ML	A212573	001	Sep 06, 2022
-----------------------------	---------	-----	--------------

ZYDUS PHARMS 0.2MG/ML	A215557	001	Apr 14, 2023
-----------------------	---------	-----	--------------

ISUPREL

+ BAUSCH 0.2MG/ML **	N010515	001
----------------------	---------	-----

SOLUTION; INHALATION

AEROLONE

LILLY 0.25%	N007245	001
-------------	---------	-----

ISOPROTERENOL HYDROCHLORIDE

ARMOUR PHARM 0.031%	A087935	001	Nov 18, 1982
---------------------	---------	-----	--------------

0.062%	A087936	001	Nov 18, 1982
--------	---------	-----	--------------

DEY 0.5%	A086764	001	Jan 04, 1982
----------	---------	-----	--------------

PARKE DAVIS 0.25%	A085994	001
-------------------	---------	-----

0.5%	A085540	001
------	---------	-----

ISUPREL

SANOFI AVENTIS US 0.5%	N006327	002
------------------------	---------	-----

1%	N006327	003
----	---------	-----

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ISOPROTERENOL HYDROCHLORIDE

SOLUTION; INHALATION

VAPO-ISO

FISONS 0.5% N016813 001

TABLET; RECTAL, SUBLINGUAL

ISUPREL

SANOFI AVENTIS US 10MG N006328 001

15MG N006328 002

ISOPROTERENOL HYDROCHLORIDE; PHENYLEPHRINE BITARTRATE

AEROSOL, METERED; INHALATION

DUO-MEDIHALER

3M 0.16MG/INH; 0.24MG/INH N013296 001

ISOPROTERENOL SULFATE

AEROSOL, METERED; INHALATION

MEDIHALER-ISO

3M 0.08MG/INH N010375 003

POWDER; INHALATION

NORISODRINE

ABBVIE 10% N006905 003

25% N006905 002

ISOSORBIDE

SOLUTION; ORAL

ISMOTIC

ALCON 100GM/220ML N017063 001

ISOSORBIDE DINITRATE

CAPSULE, EXTENDED RELEASE; ORAL

DILATRATE-SR

+ ENDO OPERATIONS 40MG N019790 001 Sep 02, 1988

ISORDIL

WYETH AYERST 40MG N012882 002 Jul 29, 1988

TABLET; ORAL

ISORDIL

+ BAUSCH 10MG ** N012093 002 Jul 29, 1988

+ 20MG ** N012093 006 Jul 29, 1988

+ 30MG ** N012093 005 Jul 29, 1988

ISOSORBIDE DINITRATE

ANI PHARMS 10MG A086032 001 Jan 07, 1988

HIKMA INTL PHARMS 30MG A040591 001 Jan 10, 2007

SUN PHARM INDUSTRIES 5MG A086166 002 Sep 19, 1986

10MG A086169 001 Sep 19, 1986

20MG A086167 001 Sep 19, 1986

30MG A087564 001 Sep 18, 1986

SUPERPHARM 5MG A089190 001 Feb 17, 1987

10MG A089191 001 Feb 17, 1987

20MG A089192 001 Feb 17, 1987

WATSON LABS 5MG A086034 001 Jan 06, 1988

SORBITRATE

ASTRAZENECA 5MG N016192 001 Apr 01, 1996

10MG N016192 002 Apr 01, 1996

20MG A086405 002 Aug 21, 1990

30MG A088124 001 Aug 21, 1990

40MG A088125 001 Aug 21, 1990

TABLET; SUBLINGUAL

ISORDIL

+ BIOVAIL 2.5MG ** N012940 004 Jul 29, 1988

+ 5MG ** N012940 003 Jul 29, 1988

+ 10MG ** N012940 005 Jul 29, 1988

ISOSORBIDE DINITRATE

HIKMA INTL PHARMS 2.5MG A086054 001 Oct 29, 1987

5MG A086055 001 Nov 02, 1987

SANDOZ 2.5MG A086225 001 Feb 19, 1988

5MG A086222 001 Feb 19, 1988

SUN PHARM INDUSTRIES 2.5MG A084204 001 Sep 18, 1986

5MG A086168 001 Sep 18, 1986

10MG A087545 001 Sep 18, 1986

WATSON LABS 2.5MG ** A086033 001 Feb 26, 1988

WATSON LABS TEVA 5MG ** A086031 001 Sep 29, 1987

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ISOSORBIDE DINITRATE

TABLET;SUBLINGUAL

SORBITRATE

ASTRAZENECA	2.5MG	N016191 002	Apr 01, 1996
	5MG	N016191 001	Apr 01, 1996

TABLET, CHEWABLE;ORAL

SORBITRATE

ASTRAZENECA	5MG	N016776 002	Apr 01, 1996
	10MG	N016776 003	Apr 01, 1996

TABLET, EXTENDED RELEASE;ORAL

ISORDIL

WYETH AYERST	40MG	N012882 001	Jul 29, 1988
--------------	------	-------------	--------------

ISOSORBIDE DINITRATE

IMPAX LABS INC	40MG	A040723 001	Mar 17, 2008
SUN PHARM INDS INC	40MG	A040009 001	Dec 30, 1998

ISOSORBIDE MONONITRATE

TABLET;ORAL

ISMO

PROMIUS PHARMA	20MG	N019091 001	Dec 30, 1991
----------------	------	-------------	--------------

ISOSORBIDE MONONITRATE

ANI PHARMS	20MG	A075147 001	Nov 27, 1998
HIKMA PHARMS	20MG	A075361 001	Oct 05, 2000

TABLET, EXTENDED RELEASE;ORAL

IMDUR

+ SCHERING PLOUGH	30MG **	N020225 001	Aug 12, 1993
	60MG **	N020225 002	Aug 12, 1993
	120MG **	N020225 003	Mar 30, 1995

ISOSORBIDE MONONITRATE

ACCORD HLTHCARE	30MG	A209684 001	Oct 24, 2017
	60MG	A209684 002	Oct 24, 2017
	120MG	A209684 003	Oct 24, 2017
ACTAVIS ELIZABETH	30MG	A075306 001	Dec 31, 1998
	60MG	A075306 002	Dec 31, 1998
ALKERMES GAINESVILLE	60MG	A075041 001	Sep 22, 1998
HIKMA INTL PHARMS	30MG	A076813 002	Mar 30, 2006
	60MG	A076813 001	Jan 07, 2005
IVAX SUB TEVA PHARMS	30MG	A075448 002	Aug 07, 2001
	60MG	A075448 001	Jun 19, 2000
	120MG	A075448 003	Aug 07, 2001
SKYEPHARMA AG	60MG	A075166 001	Oct 07, 1999
STRIDES PHARMA	30MG	A090598 001	Aug 11, 2010
	60MG	A090598 002	Aug 11, 2010
	120MG	A090598 003	Aug 11, 2010
ZYDUS PHARMS	30MG	A218255 001	Feb 15, 2024
	60MG	A218255 002	Feb 15, 2024
	120MG	A218255 003	Feb 15, 2024

ISOSULFAN BLUE

SOLUTION;SUBCUTANEOUS

ISOSULFAN BLUE

BELOTECA	50MG/5ML (10MG/ML)	A210714 001	Jan 16, 2019
EUGIA PHARMA	50MG/5ML (10MG/ML)	A206831 001	Feb 02, 2016
FRESENIUS KABI USA	50MG/5ML (10MG/ML)	A211869 001	Sep 08, 2022
SOMERSET THERAPS LLC	50MG/5ML (10MG/ML)	A210558 001	Jul 12, 2019

LYMPHAZURIN

+ COVIDIEN	50MG/5ML (10MG/ML) **	N018310 001	
------------	-----------------------	-------------	--

ISOTRETINOIN

CAPSULE;ORAL

ABSORICA LD

+ SUN PHARM	20MG	N211913 003	Nov 05, 2019
	28MG	N211913 005	Nov 05, 2019

ACCUATANE

+ HOFFMANN LA ROCHE	10MG **	N018662 002	May 07, 1982
	20MG **	N018662 004	Mar 28, 1983
	40MG **	N018662 003	May 07, 1982

SOTRET

SUN PHARM INDS LTD	10MG	A076041 001	Dec 24, 2002
	20MG	A076041 002	Dec 24, 2002
	30MG	A076503 001	Jun 20, 2003

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ISOTRETINOINCAPSULE; ORAL
SOTRET

40MG

A076041 003 Dec 24, 2002

ISRADIPINECAPSULE; ORAL
DYNACIRC

+ SMITHKLINE BEECHAM 2.5MG **

N019546 001 Dec 20, 1990

+ 5MG **

N019546 002 Dec 20, 1990

TABLET, EXTENDED RELEASE; ORAL

DYNACIRC CR

+ GLAXOSMITHKLINE LLC 5MG **

N020336 001 Jun 01, 1994

+ 10MG **

N020336 002 Jun 01, 1994

ISRADIPINE

NORVIUM BIOSCIENCE 5MG

A201067 001 Nov 27, 2015

10MG

A201067 002 Nov 27, 2015

ITRACONAZOLE

CAPSULE; ORAL

ITRACONAZOLE

ACCORD HLTHCARE 100MG

A205991 001 May 26, 2016

JUBILANT GENERICS 100MG

A203445 001 Feb 23, 2017

MYLAN PHARMS INC 100MG

A200463 001 Jul 20, 2012

RISING 100MG

A205724 001 Dec 13, 2016

STRIDES PHARMA 100MG

A206410 001 Jul 02, 2019

INJECTABLE; INJECTION

SPORANOX

JANSSEN PHARMS 10MG/ML

N020966 001 Mar 30, 1999

SOLUTION; ORAL

ITRACONAZOLE

APOTEX 10MG/ML

A208481 001 Aug 02, 2019

TABLET; ORAL

ONMEL

+ SEBELA IRELAND LTD 200MG

N022484 001 Apr 29, 2010

IVERMECTIN

LOTION; TOPICAL

IVERMECTIN

TEVA PHARMS USA 0.5%

A212485 001 Mar 21, 2022

TABLET; ORAL

STROMEKTOL

+ MERCK SHARP DOHME 6MG **

N050742 001 Nov 22, 1996

KANAMYCIN SULFATE

CAPSULE; ORAL

KANTREX

APOTHECON EQ 500MG BASE

A060516 001

EQ 500MG BASE

A061911 001

EQ 500MG BASE

A062726 001 Mar 06, 1987

INJECTABLE; INJECTION

KANAMYCIN

HIKMA EQ 75MG BASE/2ML

A062324 001

EQ 500MG BASE/2ML

A062324 002

EQ 1GM BASE/3ML

A062324 003

KANAMYCIN SULFATE

ABRAXIS PHARM EQ 75MG BASE/2ML

A062504 001 Apr 05, 1984

EQ 500MG BASE/2ML

A062504 002 Apr 05, 1984

EQ 1GM BASE/3ML

A062504 003 Apr 05, 1984

FRESENIUS KABI USA EQ 500MG BASE/2ML

A065111 001 Dec 17, 2002

EQ 1GM BASE/3ML

A065111 002 Dec 17, 2002

INTL MEDICATION EQ 500MG BASE/2ML

A062466 001 Sep 30, 1983

EQ 1GM BASE/3ML

A062466 002 Sep 30, 1983

LOCH EQ 75MG BASE/2ML

A063021 001 Jul 31, 1992

EQ 500MG BASE/2ML

A063022 001 Jul 31, 1992

EQ 1GM BASE/3ML

A063025 001 Jul 31, 1992

PHARMAFAIR EQ 75MG BASE/2ML

A062668 001 May 07, 1987

EQ 500MG BASE/2ML

A062672 001 May 07, 1987

EQ 1GM BASE/3ML

A062669 001 May 07, 1987

SOLOPAK EQ 75MG BASE/2ML

A062605 003 Feb 26, 1986

EQ 500MG BASE/2ML

A062605 001 Feb 26, 1986

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

KANAMYCIN SULFATE

INJECTABLE; INJECTION

KANAMYCIN SULFATE

	EQ 1GM BASE/3ML		A062605 002	Feb 26, 1986
WARNER CHILCOTT	EQ 1GM BASE/3ML		A063092 001	Oct 11, 1989
WATSON LABS	EQ 1GM BASE/3ML		A062520 003	May 09, 1985
KANTREX				
APOTHECON	EQ 75MG BASE/2ML		A061655 003	
	EQ 75MG BASE/2ML		A061901 003	
	EQ 75MG BASE/2ML		A062564 001	Sep 21, 1984
	EQ 500MG BASE/2ML		A061655 001	
	EQ 500MG BASE/2ML		A061901 001	
	EQ 500MG BASE/2ML		A062564 002	Sep 21, 1984
	EQ 1GM BASE/3ML		A061655 002	
	EQ 1GM BASE/3ML		A061901 002	
	EQ 1GM BASE/3ML		A062564 003	Sep 21, 1984
KLEBCIL				
KING PHARMS	EQ 75MG BASE/2ML		A062170 001	
	EQ 500MG BASE/2ML		A062170 002	
	EQ 1GM BASE/3ML		A062170 003	

KETOCONAZOLE

AEROSOL, FOAM; TOPICAL

EXTINA

+ NORVIUM BIOSCIENCE

2% **

N021738 001 Jun 12, 2007

CREAM; TOPICAL

NIZORAL

+ JANSSEN PHARMA

2% **

N019084 001 Dec 31, 1985

GEL; TOPICAL

XOLEGEL

+ ALMIRALL

2%

N021946 001 Jul 28, 2006

SHAMPOO; TOPICAL

NIZORAL

+ JANSSEN PHARMS

2% **

N019927 001 Aug 31, 1990

SUSPENSION; ORAL

NIZORAL

JANSSEN PHARMA

100MG/5ML

A070767 001 Nov 07, 1986

TABLET; ORAL

KETOCONAZOLE

AAIPHARMA LLC

200MG

A075341 001 Jul 27, 1999

AUROBINDO PHARMA USA

200MG

A075597 001 Dec 23, 1999

CHARTWELL RX

200MG

A074971 001 Jun 15, 1999

HERITAGE PHARMA

200MG

A075362 001 Jun 15, 1999

SUN PHARM INDUSTRIES

200MG

A075314 001 Jun 15, 1999

TEVA

200MG

A075273 001 Jun 15, 1999

NIZORAL

+ JANSSEN PHARMS

200MG **

N018533 001

KETOPROFEN

CAPSULE; ORAL

KETOPROFEN

RISING

50MG

A074024 001 Dec 29, 1995

50MG

A074035 002 Dec 31, 1996

75MG

A074024 002 Dec 29, 1995

75MG

A074035 003 Dec 31, 1996

TEVA

25MG

A073515 001 Dec 22, 1992

75MG

A073517 001 Dec 22, 1992

ORUDIS

+ WYETH AYERST

25MG **

N018754 001 Jul 31, 1987

+

50MG **

N018754 002 Jan 09, 1986

+

75MG **

N018754 003 Jan 09, 1986

CAPSULE, EXTENDED RELEASE; ORAL

KETOPROFEN

ACTAVIS LABS FL INC

100MG

A075270 002 Mar 24, 1999

150MG

A075270 003 Mar 24, 1999

200MG

A075270 001 Mar 24, 1999

ALKERMES GAINESVILLE

200MG

A074879 001 Dec 10, 1997

MYLAN

100MG

A075679 003 Feb 20, 2002

150MG

A075679 002 Feb 20, 2002

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

KETOPROFEN

CAPSULE, EXTENDED RELEASE;ORAL

ORUVAIL

+	WYETH PHARMS INC	100MG **	N019816	003	Feb 08, 1995
+		150MG **	N019816	002	Feb 08, 1995
+		200MG **	N019816	001	Sep 24, 1993

FILM;ORAL

NEXCEDE

	NOVARTIS	12.5MG	N022470	001	Nov 25, 2009
--	----------	--------	---------	-----	--------------

TABLET;ORAL

ACTRON

	BAYER	12.5MG	N020499	001	Oct 06, 1995
--	-------	--------	---------	-----	--------------

KETOPROFEN

	PERRIGO	12.5MG	A075364	001	Feb 07, 2002
--	---------	--------	---------	-----	--------------

ORUDIS KT

+	WYETH CONS	12.5MG **	N020429	001	Oct 06, 1995
---	------------	-----------	---------	-----	--------------

KETOROLAC TROMETHAMINE

INJECTABLE;INJECTION

KETOROLAC TROMETHAMINE

	AMPHASTAR PHARM	15MG/ML	A076209	001	Jul 21, 2004
--	-----------------	---------	---------	-----	--------------

		30MG/ML	A076209	002	Jul 21, 2004
--	--	---------	---------	-----	--------------

	APOTEX INC	30MG/ML	A075626	001	Jul 24, 2001
--	------------	---------	---------	-----	--------------

		30MG/ML	A077201	001	Oct 14, 2005
--	--	---------	---------	-----	--------------

	APOTHECON	15MG/ML	A075348	001	Nov 28, 2000
--	-----------	---------	---------	-----	--------------

		30MG/ML	A075348	002	Nov 28, 2000
--	--	---------	---------	-----	--------------

	BAXTER HLTHCARE CORP	15MG/ML	A075631	002	Jun 29, 2001
--	----------------------	---------	---------	-----	--------------

		30MG/ML	A075631	001	Jun 29, 2001
--	--	---------	---------	-----	--------------

	BEDFORD	15MG/ML	A075230	002	Oct 25, 1999
--	---------	---------	---------	-----	--------------

		30MG/ML	A075230	001	Oct 25, 1999
--	--	---------	---------	-----	--------------

	EUGIA PHARMA	15MG/ML	A212939	001	Oct 20, 2020
--	--------------	---------	---------	-----	--------------

		30MG/ML	A212939	002	Oct 20, 2020
--	--	---------	---------	-----	--------------

	GLAND PHARMA LTD	15MG/ML	A076722	001	Jul 27, 2004
--	------------------	---------	---------	-----	--------------

		30MG/ML	A076722	002	Jul 27, 2004
--	--	---------	---------	-----	--------------

	HIKMA	15MG/ML **	A075222	001	Apr 26, 1999
--	-------	------------	---------	-----	--------------

		15MG/ML	A075299	001	Nov 03, 1999
--	--	---------	---------	-----	--------------

		30MG/ML **	A075222	002	Apr 26, 1999
--	--	------------	---------	-----	--------------

		30MG/ML **	A075228	001	Apr 26, 1999
--	--	------------	---------	-----	--------------

		30MG/ML	A075299	002	Nov 03, 1999
--	--	---------	---------	-----	--------------

	HOSPIRA	15MG/ML	A074801	001	Jun 05, 1997
--	---------	---------	---------	-----	--------------

		15MG/ML	A074993	001	Jan 27, 1999
--	--	---------	---------	-----	--------------

		30MG/ML	A074801	002	Jun 05, 1997
--	--	---------	---------	-----	--------------

	LUITPOLD	15MG/ML	A078145	001	Jan 14, 2008
--	----------	---------	---------	-----	--------------

		30MG/ML	A078145	002	Jan 14, 2008
--	--	---------	---------	-----	--------------

	SANDOZ	15MG/ML	A076271	001	Oct 06, 2004
--	--------	---------	---------	-----	--------------

	STERISCIENCE SPECLTS	15MG/ML	A078299	001	Jul 16, 2007
--	----------------------	---------	---------	-----	--------------

		15MG/ML	A201155	001	Aug 04, 2014
--	--	---------	---------	-----	--------------

		30MG/ML	A078299	002	Jul 16, 2007
--	--	---------	---------	-----	--------------

		30MG/ML	A201155	002	Aug 04, 2014
--	--	---------	---------	-----	--------------

	WOCKHARDT BIO AG	30MG/ML	A077943	001	Mar 27, 2007
--	------------------	---------	---------	-----	--------------

TORADOL

+	ROCHE PALO	15MG/ML **	N019698	001	Nov 30, 1989
---	------------	------------	---------	-----	--------------

+		30MG/ML **	N019698	002	Nov 30, 1989
---	--	------------	---------	-----	--------------

SOLUTION/DROPS;OPHTHALMIC

ACULAR PRESERVATIVE FREE

	ALLERGAN	0.5%	N020811	001	Nov 03, 1997
--	----------	------	---------	-----	--------------

KETOROLAC TROMETHAMINE

	CHARTWELL RX	0.5%	A078434	001	Nov 05, 2009
--	--------------	------	---------	-----	--------------

	EPIC PHARMA LLC	0.4%	A078399	001	Nov 05, 2009
--	-----------------	------	---------	-----	--------------

		0.45%	A203376	001	Feb 10, 2014
--	--	-------	---------	-----	--------------

	EUGIA PHARMA	0.4%	A205191	001	Nov 15, 2018
--	--------------	------	---------	-----	--------------

		0.5%	A205190	001	Dec 03, 2020
--	--	------	---------	-----	--------------

	SANDOZ	0.4%	A078721	001	Nov 05, 2009
--	--------	------	---------	-----	--------------

	SUN PHARM	0.5%	A090017	001	Nov 05, 2009
--	-----------	------	---------	-----	--------------

TABLET;ORAL

KETOROLAC TROMETHAMINE

	CHARTWELL RX	10MG	A074790	001	Jun 26, 1997
--	--------------	------	---------	-----	--------------

	PLIVA	10MG	A075284	001	Jun 23, 1999
--	-------	------	---------	-----	--------------

	WATSON LABS	10MG	A074955	001	Sep 19, 1997
--	-------------	------	---------	-----	--------------

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

KETOROLAC TROMETHAMINE

TABLET; ORAL

TORADOL

+ ROCHE PALO 10MG ** N019645 001 Dec 20, 1991

KETOTIFEN FUMARATE

DRUG-ELUTING CONTACT LENS; OPHTHALMIC

ACUVUE THERAVISION WITH KETOTIFEN

+ JOHNSON JOHNSON VISN EQ 19MCG BASE N022388 001 Feb 25, 2022

SOLUTION/DROPS; OPHTHALMIC

ALAWAY

BAUSCH AND LOMB INC EQ 0.025% BASE A208158 001 Sep 24, 2020

ZADITOR

+ ALCON PHARMA EQ 0.025% BASE ** N021066 002 Oct 19, 2006

KRYPTON, KR-81M

GAS; INHALATION

MPI KRYPTON 81M GENERATOR

GE HEALTHCARE N/A N018088 001

L-GLUTAMINE

FOR SOLUTION; ORAL

NUTRESTORE

+ EMMAUS MEDCL 5GM/PACKET N021667 001 Jun 10, 2004

LABETALOL HYDROCHLORIDE

INJECTABLE; INJECTION

LABETALOL HYDROCHLORIDE

APOTHECON 5MG/ML A075355 001 Nov 29, 1999

HOSPIRA 5MG/ML A075242 001 Sep 30, 1999

RISING 5MG/ML A075524 001 Nov 29, 1999

STERISCIENCE SPECLTS 5MG/ML A079134 001 Feb 03, 2010

NORMODYNE

+ SCHERING 5MG/ML ** N018686 001 Aug 01, 1984

TRANDATE

+ SEBELA IRELAND LTD 5MG/ML ** N019425 001 Dec 31, 1985

SOLUTION; INTRAVENOUS

LABETALOL HYDROCHLORIDE

+ HIKMA 10MG/2ML (5MG/ML) N213330 005 Mar 18, 2022

TABLET; ORAL

LABETALOL HYDROCHLORIDE

APOTHECON 100MG A075223 001 Nov 20, 1998

200MG A075223 002 Nov 20, 1998

300MG A075223 003 Nov 20, 1998

TEVA 100MG A074989 001 Sep 30, 1998

200MG A074989 002 Sep 30, 1998

300MG A074989 003 Sep 30, 1998

UNICHEM 100MG A212719 001 Aug 08, 2022

200MG A212719 002 Aug 08, 2022

300MG A212719 003 Aug 08, 2022

NORMODYNE

+ SCHERING 100MG ** N018687 001 Aug 31, 1987

+ 200MG ** N018687 002 Aug 01, 1984

+ 300MG ** N018687 003 Aug 01, 1984

+ 400MG ** N018687 004 Aug 01, 1984

TRANDATE

+ ALVOGEN 100MG ** N018716 001 May 24, 1985

+ 200MG ** N018716 002 Aug 01, 1984

+ 300MG ** N018716 003 Aug 01, 1984

+ 400MG ** N018716 004 Aug 01, 1984

LACOSAMIDE

SOLUTION; ORAL

LACOSAMIDE

AMNEAL PHARMS 10MG/ML A204839 001 Mar 27, 2024

TABLET; ORAL

LACOSAMIDE

ACCORD HLTHCARE 50MG A205011 001 Jul 12, 2022

100MG A205011 002 Jul 12, 2022

150MG A205011 003 Jul 12, 2022

200MG A205011 004 Jul 12, 2022

ACTAVIS LABS FL INC 50MG A204855 001 Jan 05, 2023

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

LACOSAMIDETABLET; ORAL
LACOSAMIDE

	100MG	A204855 002	Jan 05, 2023
	150MG	A204855 003	Jan 05, 2023
	200MG	A204855 004	Jan 05, 2023
AMNEAL PHARMS	50MG	A204857 001	Mar 17, 2022
	100MG	A204857 002	Mar 17, 2022
	150MG	A204857 003	Mar 17, 2022
	200MG	A204857 004	Mar 17, 2022
APOTEX	50MG	A214567 001	Sep 23, 2022
	100MG	A214567 002	Sep 23, 2022
	150MG	A214567 003	Sep 23, 2022
	200MG	A214567 004	Sep 23, 2022
ZYDUS PHARMS	50MG	A204947 001	Dec 15, 2023
	100MG	A204947 002	Dec 15, 2023
	150MG	A204947 003	Dec 15, 2023
	200MG	A204947 004	Dec 15, 2023

LACTITOL

FOR SOLUTION; ORAL

PIZENSY

+ BRAINTREE LABS 10GM N211281 001 Feb 12, 2020

LACTULOSE

SOLUTION; ORAL

CHRONULAC

+ SANOFI AVENTIS US 10GM/15ML ** N017884 001

CONSTILAC

ALRA 10GM/15ML A071054 001 Jul 26, 1988

CONSTULOSE

ACTAVIS MID ATLANTIC 10GM/15ML A070288 001 Aug 15, 1988

DUPHALAC

SOLVAY 10GM/15ML A072372 001 Mar 22, 1989

EVALOSE

TEVA PHARMS 10GM/15ML A073497 001 May 28, 1993

LACTULOSE

ANI PHARMS 10GM/15ML A078430 001 Nov 28, 2007

HIKMA 10GM/15ML A073591 001 May 29, 1992

10GM/15ML A074076 001 Jul 03, 1995

MORTON GROVE 10GM/15ML A071841 001 Sep 22, 1988

PACO 10GM/15ML A073160 001 Aug 25, 1992

VISTAPHARM 10GM/15ML A074138 001 Sep 30, 1992

LAXILOSE

NOSTRUM LABS 10GM/15ML A073686 001 May 28, 1993

SOLUTION; ORAL, RECTAL

ACILAC

NOSTRUM LABS 10GM/15ML A073685 001 May 28, 1993

CEPHULAC

+ SANOFI AVENTIS US 10GM/15ML ** N017657 001

CHOLAC

ALRA 10GM/15ML A071331 001 Jul 26, 1988

ENULOSE

ACTAVIS MID ATLANTIC 10GM/15ML A071548 001 Aug 15, 1988

GENERLAC

MORTON GROVE 10GM/15ML A071842 001 Sep 27, 1988

HEPTALAC

TEVA PHARMS 10GM/15ML A073504 001 May 28, 1993

LACTULOSE

ANI PHARMS 10GM/15ML A090426 001 Nov 21, 2008

PACO 10GM/15ML A072029 001 Aug 25, 1992

PAI HOLDINGS PHARM 10GM/15ML A074077 001 Jul 03, 1995

ROXANE 10GM/15ML A073590 001 May 29, 1992

SOLVAY 10GM/15ML N017906 001

PORTALAC

SOLVAY 10GM/15ML A072374 001 Mar 22, 1989

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

LAMIVUDINE

SOLUTION; ORAL

EPIVIR-HBV

+ GLAXOSMITHKLINE 5MG/ML N021004 001 Dec 08, 1998

TABLET; ORAL

EPIVIR-HBV

+ GLAXOSMITHKLINE 100MG N021003 001 Dec 08, 1998

LAMIVUDINE

AUROBINDO PHARMA LTD 150MG A202032 001 Nov 17, 2011

300MG A202032 002 Nov 17, 2011

AUROBINDO PHARMA USA 100MG A204002 001 Dec 31, 2014

MYLAN LABS LTD 150MG A078545 001 Mar 05, 2019

300MG A078545 002 Mar 05, 2019

NATCO 150MG A204528 001 Mar 04, 2016

300MG A204528 002 Mar 04, 2016

LAMIVUDINE; NEVIRAPINE; ZIDOVUDINE

TABLET; ORAL

LAMIVUDINE, NEVIRAPINE AND ZIDOVUDINE

+ MICRO LABS 150MG;200MG;300MG N205626 001 Aug 13, 2018

LAMIVUDINE; RALTEGRAVIR POTASSIUM

TABLET; ORAL

DUTREBIS

MERCK SHARP DOHME 150MG;EQ 300MG BASE N206510 001 Feb 06, 2015

LAMIVUDINE; TENOFOVIR DISOPROXIL FUMARATE

TABLET; ORAL

LAMIVUDINE AND TENOFOVIR DISOPROXIL FUMARATE

+ AUROBINDO PHARMA LTD 300MG;300MG N022344 001 May 15, 2018

TEMIXYS

+ CHARTWELL RX 300MG;300MG N211284 001 Nov 16, 2018

LAMIVUDINE; ZIDOVUDINE

TABLET; ORAL

COMBIVIR

+ VIIV HLTHCARE 150MG;300MG ** N020857 001 Sep 26, 1997

LAMIVUDINE AND ZIDOVUDINE

AUROBINDO PHARMA LTD 150MG;300MG A202418 001 May 15, 2012

CHARTWELL RX 150MG;300MG A079081 001 May 25, 2011

NATCO 150MG;300MG A204005 001 Aug 28, 2014

NORVIUM BIOSCIENCE 150MG;300MG A079079 001 Aug 12, 2019

PHARMACARE 150MG;300MG N022018 001 Mar 17, 2017

LAMOTRIGINE

TABLET; ORAL

LAMICTAL

+ GLAXOSMITHKLINE LLC 50MG ** N020241 006 Dec 27, 1994

+ 250MG ** N020241 004 Dec 27, 1994

LAMOTRIGINE

ACTAVIS TOTOWA 25MG A078669 001 Apr 08, 2011

100MG A078669 002 Apr 08, 2011

150MG A078669 003 Apr 08, 2011

200MG A078669 004 Apr 08, 2011

CHARTWELL MOLECULAR 25MG A077783 001 Nov 01, 2010

100MG A077783 002 Nov 01, 2010

150MG A077783 003 Nov 01, 2010

200MG A077783 004 Nov 01, 2010

GRANULES 25MG A078982 001 Jan 27, 2009

100MG A078982 002 Jan 27, 2009

150MG A078982 003 Jan 27, 2009

200MG A078982 004 Jan 27, 2009

HIKMA PHARMS 25MG A078134 001 Apr 19, 2011

100MG A078134 002 Apr 19, 2011

150MG A078134 003 Apr 19, 2011

200MG A078134 004 Apr 19, 2011

NATCO PHARMA 25MG A077428 001 Jan 27, 2009

100MG A077428 002 Jan 27, 2009

150MG A077428 003 Jan 27, 2009

200MG A077428 004 Jan 27, 2009

NORVIUM BIOSCIENCE 25MG A078443 001 Feb 11, 2009

100MG A078443 002 Feb 11, 2009

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

LAMOTRIGINE

TABLET;ORAL

LAMOTRIGINE

	150MG	A078443 003	Feb 11, 2009
	200MG	A078443 004	Feb 11, 2009
RISING	25MG	A077420 001	Jan 27, 2009
	100MG	A077420 002	Jan 27, 2009
	150MG	A077420 003	Jan 27, 2009
	200MG	A077420 004	Jan 27, 2009
ROXANE	25MG	A077392 001	Jan 27, 2009
	100MG	A077392 002	Jan 27, 2009
	150MG	A077392 003	Jan 27, 2009
	200MG	A077392 004	Jan 27, 2009
SANDOZ	25MG	A078645 001	Jan 27, 2009
	100MG	A078645 002	Jan 27, 2009
	150MG	A078645 003	Jan 27, 2009
	200MG	A078645 004	Jan 27, 2009
TEVA	25MG	A076388 001	Aug 30, 2006
	100MG	A076388 002	Aug 30, 2006
	150MG	A076388 003	Aug 30, 2006
	200MG	A076388 004	Aug 30, 2006
ZENNOVA	25MG	A078310 001	Feb 04, 2009
	100MG	A078310 002	Feb 04, 2009
	150MG	A078310 003	Feb 04, 2009
	200MG	A078310 004	Feb 04, 2009

TABLET, EXTENDED RELEASE;ORAL

LAMOTRIGINE

ENDO OPERATIONS	25MG	A201374 001	Dec 26, 2012
	50MG	A201374 002	Dec 26, 2012
	100MG	A201374 003	Dec 26, 2012
	200MG	A201374 004	Dec 26, 2012
	250MG	A201374 005	Dec 26, 2012
	300MG	A201374 006	Dec 26, 2012
RUBICON	25MG	A202887 001	Jun 17, 2013
	50MG	A202887 002	Jun 17, 2013
TORRENT	200MG	A203370 004	Dec 23, 2013

TABLET, FOR SUSPENSION;ORAL

LAMICTAL CD

GLAXOSMITHKLINE LLC	100MG	N020764 003	Aug 24, 1998
---------------------	-------	-------------	--------------

LAMOTRIGINE

CHARTWELL RX	5MG	A078409 002	Jan 22, 2009
	25MG	A078409 003	Jan 22, 2009
JUBILANT GENERICS	5MG	A200220 001	Feb 28, 2011
	25MG	A200220 002	Feb 28, 2011
NORVIUM BIOSCIENCE	5MG	A076630 001	Jan 22, 2009
	25MG	A076630 002	Jan 22, 2009
TEVA	5MG	A076420 001	Jun 21, 2006
	25MG	A076420 002	Jun 21, 2006

LANSOPRAZOLE

CAPSULE, DELAYED REL PELLETS;ORAL

LANSOPRAZOLE

AJANTA PHARMA LTD	15MG	A203957 001	Oct 14, 2016
	30MG	A203957 002	Oct 14, 2016
ESJAY PHARMA	15MG	A203964 001	Oct 17, 2018
	30MG	A203964 002	Oct 17, 2018
KRKA TOVARNA ZDRAVIL	15MG	A091212 001	Sep 16, 2013
	30MG	A091212 002	Sep 16, 2013
LANNETT CO INC	15MG	A207157 001	Sep 29, 2017
MYLAN	15MG	A203187 001	Jun 01, 2016
SANDOZ	15MG	A090331 001	Apr 23, 2010
	30MG	A090331 002	Apr 23, 2010
SUN PHARM	15MG	A202637 001	Sep 13, 2013
	30MG	A091509 001	Sep 13, 2013

PREVACID

+ TAKEDA PHARMS USA 15MG **

N020406 001 May 10, 1995

FOR SUSPENSION, DELAYED RELEASE;ORAL

PREVACID

TAKEDA PHARMS NA	15MG/PACKET	N021281 001	May 03, 2001
	30MG/PACKET	N021281 002	May 03, 2001

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

LANSOPRAZOLE

INJECTABLE; INTRAVENOUS

PREVACID IV

+ TAKEDA PHARMS NA 30MG/VIAL **

N021566 001 May 27, 2004

TABLET, ORALLY DISINTEGRATING, DELAYED RELEASE; ORAL

LANSOPRAZOLE

ANI PHARMS

15MG

A078730 001 Oct 15, 2010

30MG

A078730 002 Oct 15, 2010

LANSOPRAZOLE; NAPROXEN

CAPSULE, DELAYED REL PELLETS, TABLET; ORAL

PREVACID NAPRAPAC 250 (COPACKAGED)

+ TAKEDA PHARMS NA 15MG, N/A; N/A, 250MG **

N021507 002 Nov 14, 2003

PREVACID NAPRAPAC 375 (COPACKAGED)

TAKEDA PHARMS NA 15MG, N/A; N/A, 375MG

N021507 003 Nov 14, 2003

PREVACID NAPRAPAC 500 (COPACKAGED)

TAKEDA PHARMS NA 15MG, N/A; N/A, 500MG

N021507 004 Nov 14, 2003

LANTHANUM CARBONATE

TABLET, CHEWABLE; ORAL

FOSRENOL

TAKEDA PHARMS USA EQ 250MG BASE

N021468 001 Oct 26, 2004

LAPYRIUM CHLORIDE; UNDECOYLIUM CHLORIDE IODINE COMPLEX

SOLUTION; TOPICAL

VIRAC REX

CHESEBROUGH PONDS 0.5%; 1.8%

N011914 001

LASMIDITAN SUCCINATE

TABLET; ORAL

REYVOW

+ ELI LILLY AND CO EQ 200MG BASE

N211280 003 Dec 18, 2020

LATANOPROST

EMULSION; OPHTHALMIC

XELPROS

+ SUN PHARM 0.005%

N206185 001 Sep 12, 2018

SOLUTION/DROPS; OPHTHALMIC

LATANOPROST

APOTEX INC 0.005%

A077697 001 Mar 22, 2011

EPIC PHARMA LLC 0.005%

A090887 001 Jul 19, 2011

EUGIA PHARMA 0.005%

A206519 001 Sep 03, 2019

RYAN LABS 0.005%

A202077 001 Feb 11, 2013

LEFLUNOMIDE

TABLET; ORAL

LEFLUNOMIDE

BARR 10MG

A077083 001 Sep 13, 2005

20MG

A077083 002 Sep 13, 2005

SANDOZ 10MG

A077085 001 Sep 13, 2005

20MG

A077085 002 Sep 13, 2005

TEVA PHARMS 10MG

A077084 001 Sep 13, 2005

20MG

A077084 002 Sep 13, 2005

LENALIDOMIDE

CAPSULE; ORAL

LENALIDOMIDE

QILU 2.5MG

A217265 001 Feb 22, 2024

5MG

A217265 002 Feb 22, 2024

10MG

A217265 003 Feb 22, 2024

15MG

A217265 004 Feb 22, 2024

20MG

A217265 005 Feb 22, 2024

25MG

A217265 006 Feb 22, 2024

TORRENT 10MG

A213405 001 Feb 17, 2023

20MG

A213405 003 Aug 03, 2023

25MG

A213405 002 Feb 17, 2023

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

LESINURAD

TABLET;ORAL

ZURAMPIC

+ IRONWOOD PHARMS INC 200MG N207988 001 Dec 22, 2015

LETROZOLE

TABLET;ORAL

LETROZOLE

ACTAVIS TOTOWA	2.5MG	A090292 001	Jul 13, 2011
APOTEX INC	2.5MG	A091303 001	Apr 19, 2012
CHARTWELL RX	2.5MG	A202716 001	May 16, 2013
FRESENIUS KABI USA	2.5MG	A090491 001	Jun 03, 2011
HIKMA	2.5MG	A090838 001	Jun 03, 2011
HIKMA PHARMS	2.5MG	A203796 001	Jun 03, 2016
IMPAX LABS	2.5MG	A091638 001	Jun 03, 2011
INDICUS PHARMA	2.5MG	A201804 001	Jun 03, 2011
LANNETT CO INC	2.5MG	A091098 001	Jun 03, 2011
	2.5MG	A202048 001	Oct 29, 2014
NORVIUM BIOSCIENCE	2.5MG	A078190 001	Dec 24, 2008
RYAN LABS	2.5MG	A091191 001	Jun 03, 2011
STRIDES PHARMA	2.5MG	A090789 001	Jun 03, 2011
SUN PHARM INDS LTD	2.5MG	A091466 001	Jun 03, 2011
SYNTHON PHARMS	2.5MG	A090196 001	Jun 03, 2011

LEUCOVORIN CALCIUM

FOR SOLUTION;ORAL

LEUCOVORIN CALCIUM

HOSPIRA EQ 60MG BASE/VIAL N008107 003 Jan 30, 1987

INJECTABLE; INJECTION

LEUCOVORIN CALCIUM

ABIC	EQ 50MG BASE/VIAL	A089353 001	Jun 01, 1988
ABRAXIS PHARM	EQ 50MG BASE/VIAL	A088939 001	Dec 01, 1986
ELKINS SINN	EQ 50MG BASE/VIAL	A070480 001	Jan 02, 1987
	EQ 100MG BASE/VIAL	A081224 001	Jun 03, 1994
+ HOSPIRA	EQ 50MG BASE/VIAL **	N008107 002	
+	EQ 100MG BASE/VIAL **	N008107 004	May 23, 1988
+	EQ 350MG BASE/VIAL **	N008107 005	Apr 05, 1989
PHARMACHEMIE	EQ 350MG BASE/VIAL	A040262 001	Dec 15, 1999
PHARMACHEMIE USA	EQ 50MG BASE/VIAL	A089628 001	Apr 17, 1997
	EQ 100MG BASE/VIAL	A089915 001	Apr 17, 1997
TEVA PARENTERAL	EQ 50MG BASE/VIAL	A081278 001	Sep 28, 1993
TEVA PHARMS USA	EQ 100MG BASE/VIAL	A081277 001	Sep 28, 1993
	EQ 350MG BASE/VIAL	A040174 001	Jun 12, 1997

LEUCOVORIN CALCIUM PRESERVATIVE FREE

AM REGENT EQ 50MG BASE/VIAL A040338 001 Jan 31, 2001

WELLCOVORIN

GLAXOSMITHKLINE	EQ 25MG BASE/VIAL	A089833 001	Jan 23, 1989
	EQ 50MG BASE/VIAL	A089465 001	Jan 23, 1989
	EQ 100MG BASE/VIAL	A089834 001	Jan 23, 1989

SOLUTION;INTRAMUSCULAR, INTRAVENOUS

LEUCOVORIN CALCIUM

ABIC	EQ 3MG BASE/ML (EQ 3MG BASE/ML)	A089352 001	Jun 01, 1988
+ HOSPIRA	EQ 3MG BASE/ML (EQ 3MG BASE/ML) **	N008107 001	
NOVAST LABS	EQ 500MG BASE/50ML (EQ 10MG BASE/ML)	A210917 001	Nov 23, 2018

LEUCOVORIN CALCIUM PRESERVATIVE FREE

HIKMA	EQ 300MG BASE/30ML (EQ 10MG BASE/ML)	A040347 001	Apr 25, 2000
	EQ 500MG BASE/50ML (EQ 10MG BASE/ML)	A040347 002	Apr 25, 2000
HOSPIRA	EQ 100MG BASE/10ML (EQ 10MG BASE/ML) **	A040147 001	Jun 25, 1997
	EQ 250MG BASE/25ML (EQ 10MG BASE/ML)	A040147 002	Jun 25, 1997
TEVA PARENTERAL	EQ 500MG BASE/50ML (EQ 10MG BASE/ML)	A040332 001	Jun 28, 1999

WELLCOVORIN

GLAXOSMITHKLINE	EQ 5MG BASE/ML (EQ 5MG BASE/ML)	A087439 001	Oct 19, 1982
	EQ 25MG BASE/5ML (EQ 5MG BASE/ML)	A087439 002	Oct 19, 1982

TABLET;ORAL

LEUCOVORIN CALCIUM

ANI PHARMS	EQ 15MG BASE	A075327 001	Mar 24, 1999
PAR PHARM	EQ 5MG BASE	A071600 001	Oct 14, 1987
	EQ 25MG BASE	A071598 001	Oct 14, 1987
PHARMACHEMIE	EQ 5MG BASE	A073099 001	Mar 28, 1997
	EQ 25MG BASE	A073101 001	Mar 28, 1997

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

LEUCOVORIN CALCIUM

TABLET; ORAL

LEUCOVORIN CALCIUM

XANODYNE PHARM

EQ 5MG BASE

N018459 001 Jan 30, 1986

EQ 10MG BASE

A071962 001 Nov 19, 1987

EQ 15MG BASE

A071104 001 Mar 04, 1987

WELLCOVORIN

+ GLAXOSMITHKLINE

EQ 5MG BASE **

N018342 001 Jul 08, 1983

+

EQ 25MG BASE **

N018342 002 Jul 08, 1983

LEUPROLIDE ACETATE

IMPLANT; IMPLANTATION

VIADUR

ORTHO MCNEIL JANSSEN EQ 65MG BASE

N021088 001 Mar 03, 2000

INJECTABLE; INJECTION

LUPRON DEPOT

+ ABBVIE ENDOCRINE INC

3.75MG/VIAL **

N020011 001 Oct 22, 1990

POWDER; INTRAMUSCULAR

LUPRON DEPOT-PED KIT

+ ABBVIE ENDOCRINE INC

3.75MG, 7.5MG **

N020263 003 Apr 16, 1993

+

7.5MG, 7.5MG **

N020263 004 Apr 16, 1993

SOLUTION; SUBCUTANEOUS

LEUPROLIDE ACETATE

GENZYME

14MG/2.8ML (1MG/0.2ML)

A075721 001 Nov 29, 2001

LUPRON

+ ABBVIE ENDOCRINE INC

14MG/2.8ML (1MG/0.2ML) **

N019010 001 Apr 09, 1985

LEUPROLIDE ACETATE; NORETHINDRONE ACETATE

INJECTABLE, TABLET; INTRAMUSCULAR, ORAL

LUPANETA PACK

+ ABBVIE ENDOCRINE

3.75MG/VIAL, N/A; N/A, 5MG

N203696 001 Dec 14, 2012

+

11.25MG/VIAL, N/A; N/A, 5MG

N203696 002 Dec 14, 2012

LEVALBUTEROL HYDROCHLORIDE

SOLUTION; INHALATION

LEVALBUTEROL HYDROCHLORIDE

NORVIUM BIOSCIENCE

EQ 0.0103% BASE

A077800 001 Mar 15, 2013

EQ 0.021% BASE

A077800 002 Mar 15, 2013

EQ 0.042% BASE

A077800 003 Mar 15, 2013

SUN PHARM

EQ 0.0103% BASE

A207820 001 Nov 05, 2018

EQ 0.021% BASE

A207820 002 Nov 05, 2018

EQ 0.042% BASE

A207820 003 Nov 05, 2018

XOPENEX

+ HIKMA

EQ 0.0103% BASE **

N020837 003 Jan 30, 2002

+

EQ 0.021% BASE **

N020837 001 Mar 25, 1999

+

EQ 0.042% BASE **

N020837 002 Mar 25, 1999

+

EQ 0.25% BASE **

N020837 004 Jul 18, 2003

LEVALLORPHAN TARTRATE

INJECTABLE; INJECTION

LORFAN

ROCHE

1MG/ML

N010423 001

LEVAMISOLE HYDROCHLORIDE

TABLET; ORAL

ERGAMISOL

JANSSEN PHARMA

EQ 50MG BASE

N020035 001 Jun 18, 1990

LEVAMLODIPINE MALEATE

TABLET; ORAL

CONJUPRI

+ CSPC OUYI

EQ 1.25MG BASE

N212895 001 Dec 19, 2019

+

EQ 2.5MG BASE

N212895 002 Dec 19, 2019

+

EQ 5MG BASE

N212895 003 Dec 19, 2019

LEVETIRACETAM

INJECTABLE; INTRAVENOUS

LEVETIRACETAM

AM REGENT

500MG/5ML (100MG/ML)

A202143 001 Jan 31, 2012

EPIC PHARMA LLC

500MG/5ML (100MG/ML)

A209934 001 May 04, 2018

FRESENIUS KABI USA

500MG/5ML (100MG/ML)

A090813 001 May 26, 2010

JUBILANT GENERICS

500MG/5ML (100MG/ML)

A206838 001 Jun 02, 2016

SUN PHARM INDS LTD

500MG/5ML (100MG/ML)

A090754 001 Jun 16, 2010

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

LEVETIRACETAM

INJECTABLE; INTRAVENOUS

LEVETIRACETAM IN SODIUM CHLORIDE

FRESENIUS KABI USA	500MG/100ML (5MG/ML)	A208619 001	Jan 31, 2023
	1GM/100ML (10MG/ML)	A208619 002	Jan 31, 2023
	1.5GM/100ML (15MG/ML)	A208619 003	Jan 31, 2023
+ HQ SPCLT PHARMA	250MG/50ML (5MG/ML) **	N202543 004	Dec 14, 2020

SOLUTION; ORAL

LEVETIRACETAM

APOTEX INC	100MG/ML	A090187 001	Aug 05, 2011
BIONPHARMA	100MG/ML	A079120 001	Jan 16, 2009
HIKMA	100MG/ML	A090601 001	Feb 28, 2012
PHARMOBEDIANT CNSLTG	100MG/ML	A090028 001	Mar 03, 2010
STRIDES PHARMA	100MG/ML	A214673 001	Nov 20, 2023
TOLMAR	100MG/ML	A079107 001	Jan 15, 2009

TABLET; ORAL

LEVETIRACETAM

ACCORD HLTHCARE	250MG	A090843 001	Feb 14, 2011
	500MG	A090843 002	Feb 14, 2011
	750MG	A090843 003	Feb 14, 2011
	1GM	A090843 004	Feb 14, 2011
ACTAVIS LABS FL INC	250MG	A077408 001	Mar 02, 2009
	500MG	A077408 002	Mar 02, 2009
	750MG	A077408 003	Mar 02, 2009
FOSUN PHARMA	250MG	A077324 001	Jan 15, 2009
	500MG	A077324 002	Jan 15, 2009
	750MG	A077324 003	Jan 15, 2009
	1GM	A077324 004	Jan 15, 2009
GRANULES	250MG	A217878 001	Jun 13, 2023
	500MG	A217878 002	Jun 13, 2023
	750MG	A217878 003	Jun 13, 2023
	1GM	A217878 004	Jun 13, 2023
INVAGEN PHARMS	250MG	A078234 001	Jan 15, 2009
	500MG	A078234 002	Jan 15, 2009
	750MG	A078234 003	Jan 15, 2009
LOTUS PHARM CO LTD	250MG	A090906 002	Oct 31, 2016
	500MG	A090906 001	Nov 05, 2010
	750MG	A090906 003	Oct 31, 2016
	1GM	A090906 004	Oct 31, 2016
MYLAN	250MG	A076919 001	Nov 04, 2008
	250MG	A078731 001	Feb 10, 2009
	500MG	A078731 002	Feb 10, 2009
	750MG	A078731 003	Feb 10, 2009
	1GM	A078731 004	Feb 10, 2009
NOSTRUM LABS INC	250MG	A090511 001	Aug 18, 2011
	500MG	A090511 002	Aug 18, 2011
	750MG	A090511 003	Aug 18, 2011
	1GM	A090511 004	Aug 18, 2011
SECAN PHARMS	1GM	A205102 003	Dec 16, 2015
	500MG	A205102 004	Dec 16, 2015
TEVA PHARMS	250MG	A078101 001	Jan 15, 2009
	500MG	A078101 002	Jan 15, 2009
	750MG	A078101 003	Jan 15, 2009
	1GM	A078101 004	Jan 15, 2009
WATSON LABS INC	250MG	A078797 002	Jan 15, 2009
	500MG	A078797 003	Jan 15, 2009
	750MG	A078797 004	Jan 15, 2009
	1GM	A078797 001	Jan 15, 2009

TABLET, EXTENDED RELEASE; ORAL

ELEPSIA XR

+ TRIPOINT	1GM	N204417 001	Dec 20, 2018
+	1.5GM	N204417 002	Dec 20, 2018

LEVETIRACETAM

ACTAVIS ELIZABETH	500MG	A091557 001	Sep 12, 2011
	750MG	A091557 002	Sep 12, 2011
ADAPTIS	500MG	A202167 001	Sep 04, 2015
	750MG	A202167 002	Sep 04, 2015
AUROBINDO PHARMA USA	500MG	A200475 001	Dec 19, 2011
	750MG	A200475 002	Dec 19, 2011

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

LEVETIRACETAMTABLET, EXTENDED RELEASE;ORAL
LEVETIRACETAM

	1GM	A200475	003	Dec 07, 2015
CHARTWELL RX	500MG	A091291	001	Sep 12, 2011
	750MG	A091291	002	Sep 12, 2011
ENDO OPERATIONS	500MG	A091360	001	Oct 04, 2011
	750MG	A091360	002	Oct 04, 2011
LOTUS PHARM CO LTD	500MG	A202095	002	Jun 06, 2016
	750MG	A202095	001	Jun 06, 2016
PHARMADAX INC	500MG	A201464	001	May 25, 2012
	750MG	A201464	002	May 25, 2012
PRINSTON INC	500MG	A202533	001	Jul 20, 2012
	750MG	A202533	002	Jul 20, 2012
ROUSES POINT PHARMS	500MG	A202524	001	Aug 27, 2012
	750MG	A202524	002	Aug 27, 2012
SANDOZ	500MG	A091668	001	Nov 01, 2012
	750MG	A091668	002	Nov 01, 2012
SUN PHARM	500MG	A203059	001	Sep 09, 2013
	750MG	A203059	002	Sep 09, 2013
SUN PHARM INDUSTRIES	500MG	A091285	001	Sep 12, 2011
	750MG	A091285	002	Sep 12, 2011
TEVA PHARMS	500MG	A091430	001	Sep 12, 2011
	750MG	A091430	002	Sep 12, 2011
TORRENT PHARMS LTD	500MG	A091338	001	May 29, 2012
	750MG	A091338	002	May 29, 2012

LEVOBETAXOLOL HYDROCHLORIDESUSPENSION/DROPS;OPHTHALMIC
BETAXON

ALCON PHARMS LTD EQ 0.5% BASE N021114 001 Feb 23, 2000

LEVOBUNOLOL HYDROCHLORIDESOLUTION/DROPS;OPHTHALMIC
AKBETAEPIC PHARMA LLC 0.25% A074779 001 Oct 29, 1996
0.5% A074780 001 Oct 29, 1996

BETAGAN

+ ALLERGAN 0.25% ** N019814 001 Jun 28, 1989

LEVOBUNOLOL HYDROCHLORIDE

ALCON LABS INC 0.25% A074851 001 Oct 28, 1996
APOTEX INC 0.25% A075473 001 Aug 03, 2000
0.5% A075475 001 Aug 03, 2000
BAUSCH AND LOMB 0.25% A074307 001 Mar 04, 1994
CHARTWELL RX 0.5% A074850 001 Oct 28, 1996LEVOBUPIVACAINE HYDROCHLORIDE

INJECTABLE; INJECTION

CHIROCAINE

+ PURDUE PHARMA LP EQ 2.5MG BASE/ML ** N020997 001 Aug 05, 1999

+ EQ 5MG BASE/ML ** N020997 002 Aug 05, 1999

+ EQ 7.5MG BASE/ML ** N020997 003 Aug 05, 1999

LEVOCABASTINE HYDROCHLORIDE

SUSPENSION/DROPS;OPHTHALMIC

LIVOSTIN

NOVARTIS EQ 0.05% BASE N020219 001 Nov 10, 1993

LEVOCARNITINE

INJECTABLE; INJECTION

LEVOCARNITINE

TEVA PHARMS USA 200MG/ML A075881 001 Mar 29, 2001

SOLUTION; ORAL

CARNITOR

LEADIANT BIOSCI INC 1GM/10ML N018948 002 Apr 27, 1988

LEVOCARNITINE

HIKMA 1GM/10ML A077399 001 Oct 25, 2007

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

LEVOCETIRIZINE DIHYDROCHLORIDE

SOLUTION; ORAL

LEVOCETIRIZINE DIHYDROCHLORIDE

APOTEX 2.5MG/5ML

A202915 001 Aug 21, 2014

XYZAL

+ CHATTEM SANOFI 2.5MG/5ML **

N022157 001 Jan 28, 2008

TABLET; ORAL

LEVOCETIRIZINE DIHYDROCHLORIDE

APOTEX 5MG

A203027 001 Feb 13, 2015

GRANULES 5MG

A090486 001 Mar 26, 2013

IPCA LABS LTD 5MG

A204626 001 May 06, 2024

SUN PHARM INDS LTD 5MG

A201653 001 Jun 26, 2015

US ANTIBIOTICS 5MG

A204323 001 Dec 20, 2016

XYZAL

+ CHATTEM SANOFI 5MG **

N022064 001 May 25, 2007

LEVODOPA

CAPSULE; ORAL

BENDOPA

VALEANT PHARM INTL 100MG

N016948 003

250MG

N016948 001

500MG

N016948 002

DOPAR

SHIRE 100MG

N016913 003

250MG

N016913 001

500MG

N016913 002

LARODOPA

ROCHE 100MG

N016912 002

250MG

N016912 001

500MG

N016912 006

TABLET; ORAL

DOPAR

SHIRE 250MG

N016913 004

500MG

N016913 005

LARODOPA

ROCHE 100MG

N016912 005

250MG

N016912 003

500MG

N016912 004

LEVOFLOXACIN

INJECTABLE; INJECTION

LEVAQUIN

+ JANSSEN PHARMS EQ 500MG/20ML (EQ 25MG/ML) **

N020635 001 Dec 20, 1996

+ EQ 750MG/30ML (EQ 25MG/ML) **

N020635 004 Dec 20, 1996

LEVAQUIN IN DEXTROSE 5% IN PLASTIC CONTAINER

+ JANSSEN PHARMS EQ 250MG/50ML (EQ 5MG/ML) **

N020635 002 Dec 20, 1996

+ EQ 500MG/100ML (EQ 5MG/ML) **

N020635 003 Dec 20, 1996

+ EQ 750MG/150ML (EQ 5MG/ML) **

N020635 005 Dec 20, 1996

LEVOFLOXACIN

AVET LIFESCIENCES EQ 500MG/20ML (EQ 25MG/ML)

A202590 001 Jan 24, 2013

EQ 750MG/30ML (EQ 25MG/ML)

A202590 002 Jan 24, 2013

BAXTER HLTHCARE CORP EQ 500MG/20ML (EQ 25MG/ML)

A091436 001 Jun 05, 2013

EUGIA PHARMA EQ 500MG/20ML (EQ 25MG/ML)

A202328 001 Jan 24, 2013

EQ 750MG/30ML (EQ 25MG/ML)

A202328 002 Jan 24, 2013

HOSPIRA EQ 500MG/20ML (EQ 25MG/ML)

A078577 001 Aug 12, 2015

EQ 750MG/30ML (EQ 25MG/ML)

A078577 002 Aug 12, 2015

NORVIUM BIOSCIENCE EQ 500MG/20ML (EQ 25MG/ML)

A200560 001 Jun 20, 2011

EQ 750MG/30ML (EQ 25MG/ML)

A200560 002 Jun 20, 2011

ZYDUS PHARMS EQ 500MG/20ML (EQ 25MG/ML)

A205968 001 Jun 01, 2017

EQ 750MG/30ML (EQ 25MG/ML)

A205968 002 Jun 01, 2017

LEVOFLOXACIN IN DEXTROSE 5% IN PLASTIC CONTAINER

BAXTER HLTHCARE CORP EQ 250MG/50ML (EQ 5MG/ML)

A091397 001 Aug 08, 2013

EQ 500MG/100ML (EQ 5MG/ML)

A091397 002 Aug 08, 2013

EQ 750MG/150ML (EQ 5MG/ML)

A091397 003 Aug 08, 2013

EUGIA PHARMA EQ 250MG/50ML (EQ 5MG/ML)

A206919 001 Feb 10, 2016

EQ 500MG/100ML (EQ 5MG/ML)

A206919 002 Feb 10, 2016

EQ 750MG/150ML (EQ 5MG/ML)

A206919 003 Feb 10, 2016

KNACK EQ 250MG/50ML (EQ 5MG/ML)

A216164 001 Jan 29, 2024

EQ 500MG/100ML (EQ 5MG/ML)

A216164 002 Jan 29, 2024

EQ 750MG/150ML (EQ 5MG/ML)

A216164 003 Jan 29, 2024

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

LEVOFLOXACIN

SOLUTION; ORAL

LEVAQUIN

+ JANSSEN PHARMS 250MG/10ML N021721 001 Oct 21, 2004

SOLUTION/DROPS; OPHTHALMIC

IQUIX

+ SANTEN 1.5% ** N021571 001 Mar 01, 2004

LEVOFLOXACIN

NORVIUM BIOSCIENCE 0.5% A204899 001 Dec 08, 2017

SCIEGEN PHARMS INC 0.5% A090268 001 Dec 20, 2010

WATSON LABS TEVA 0.5% A076826 001 Feb 10, 2011

QUIXIN

+ SANTEN 0.5% ** N021199 001 Aug 18, 2000

TABLET; ORAL

LEVAQUIN

+ JANSSEN PHARMS 250MG ** N020634 001 Dec 20, 1996

+ 500MG ** N020634 002 Dec 20, 1996

+ 750MG ** N020634 003 Sep 08, 2000

LEVOFLOXACIN

ANDA REPOSITORY 250MG A090367 001 Jun 20, 2011

500MG A090367 002 Jun 20, 2011

750MG A090367 003 Jun 20, 2011

JUBILANT GENERICS 250MG A203613 001 Jun 19, 2015

500MG A203613 002 Jun 19, 2015

250MG A076276 001 Jun 20, 2011

500MG A076276 002 Jun 20, 2011

750MG A077097 001 Jun 20, 2011

SANDOZ 250MG A077438 001 Jun 20, 2011

500MG A077438 002 Jun 20, 2011

750MG A077438 003 Jun 20, 2011

TORRENT PHARMS 250MG A090722 001 Jun 20, 2011

500MG A090722 002 Jun 20, 2011

750MG A090722 003 Jun 20, 2011

WATSON LABS INC 250MG A201484 001 Nov 22, 2013

500MG A201484 002 Nov 22, 2013

750MG A201484 003 Nov 22, 2013

LEVOLEUCOVORIN

POWDER; INTRAVENOUS

KHAPZORY

+ ACROTECH BIOPHARMA 300MG/VIAL N211226 002 Oct 19, 2018

LEVOLEUCOVORIN CALCIUM

POWDER; INTRAVENOUS

FUSILEV

+ ACROTECH BIOPHARMA EQ 50MG BASE/VIAL N020140 001 Mar 07, 2008

LEVOLEUCOVORIN CALCIUM

ACTAVIS LLC EQ 50MG BASE/VIAL A206516 001 Feb 13, 2017

+ EQ 175MG BASE/VIAL N208723 001 Sep 29, 2016

AMNEAL EQ 50MG BASE/VIAL A207547 001 Feb 13, 2017

HIKMA EQ 50MG BASE/VIAL A206263 001 Jun 16, 2016

MEITHEAL EQ 50MG BASE/VIAL A211003 001 Aug 22, 2019

SOLUTION; INTRAVENOUS

FUSILEV

+ ACROTECH BIOPHARMA EQ 175MG BASE/17.5ML (EQ 10MG BASE/ML) N020140 002 Apr 29, 2011

+ EQ 250MG BASE/25ML (EQ 10MG BASE/ML) ** N020140 003 Apr 29, 2011

LEVOLEUCOVORIN CALCIUM

MEITHEAL EQ 250MG BASE/25ML (EQ 10MG BASE/ML) A211002 002 Aug 16, 2019

NORVIUM BIOSCIENCE EQ 175MG BASE/17.5ML (EQ 10MG BASE/ML) A203576 001 Oct 20, 2015

EQ 250MG BASE/25ML (EQ 10MG BASE/ML) A203576 002 Oct 20, 2015

NOVAST LABS EQ 175MG BASE/17.5ML (EQ 10MG BASE/ML) A210623 001 May 03, 2018

EQ 250MG BASE/25ML (EQ 10MG BASE/ML) A210623 002 May 03, 2018

PRAXGEN EQ 175MG BASE/17.5ML (EQ 10MG BASE/ML) A213797 001 Nov 02, 2021

SANDOZ EQ 250MG BASE/25ML (EQ 10MG BASE/ML) A203563 002 Mar 09, 2015

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

LEVOMEPRMAZINE

INJECTABLE; INJECTION

LEVOPROME

IMMUNEX

20MG/ML

N015865 001

LEVOMETHADYL ACETATE HYDROCHLORIDE

CONCENTRATE; ORAL

ORLAAM

+ ROXANE

10MG/ML **

N020315 001 Jul 09, 1993

LEVOMILNACIPRAN HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

LEVOMILNACIPRAN HYDROCHLORIDE

AMNEAL PHARMS CO

EQ 20MG BASE

A210790 001 Feb 04, 2019

EQ 40MG BASE

A210790 002 Feb 04, 2019

EQ 80MG BASE

A210790 003 Feb 04, 2019

EQ 120MG BASE

A210790 004 Feb 04, 2019

AUROBINDO PHARMA LTD

EQ 20MG BASE

A210826 001 Jan 06, 2023

EQ 40MG BASE

A210826 002 Jan 06, 2023

EQ 80MG BASE

A210826 003 Jan 06, 2023

EQ 120MG BASE

A210826 004 Jan 06, 2023

HIKMA

EQ 20MG BASE

A210732 001 Nov 05, 2020

EQ 40MG BASE

A210732 002 Nov 05, 2020

EQ 80MG BASE

A210732 003 Nov 05, 2020

EQ 120MG BASE

A210732 004 Nov 05, 2020

LEVONORDEFIN; MEPIVACAINE HYDROCHLORIDE

INJECTABLE; INJECTION

ARESTOCAINE HYDROCHLORIDE W/ LEVONORDEFIN

SOLVAY

0.05MG/ML; 2%

A085010 001

CARBOCAINE W/ NEO-COBEFRIN

EASTMAN KODAK

0.05MG/ML; 2%

N012125 002

ISOCAINE HYDROCHLORIDE W/ LEVONORDEFIN

SEPTODONT

0.05MG/ML; 2%

A084697 001

MEPIVACAINE HYDROCHLORIDE W/ LEVONORDEFIN

BELMORA LLC

0.05MG/ML; 2%

A084850 002 Oct 21, 1983

POLOCAINE W/ LEVONORDEFIN

DENTSPLY PHARM

0.05MG/ML; 2%

A089517 001 Apr 14, 1988

SCANDONEST L

DEPROCO

0.05MG/ML; 2%

A088388 001 Oct 10, 1984

LEVONORDEFIN; PROCAINE HYDROCHLORIDE; PROPOXYCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

RAVOCAINE AND NOVOCAIN W/ NEO-COBEFRIN

EASTMAN KODAK

0.05MG/ML; 2%; 0.4%

N008592 007

LEVONORGESTREL

IMPLANT; IMPLANTATION

JADELLE

+ POPULATION COUNCIL

75MG/IMPLANT **

N020544 001 Nov 01, 1996

LEVONORGESTREL

WYETH PHARMS INC

75MG/IMPLANT

N020627 001 Aug 15, 1996

NORPLANT

POPULATION COUNCIL

36MG/IMPLANT

N019897 001 Dec 10, 1990

NORPLANT SYSTEM IN PLASTIC CONTAINER

WYETH PHARMS INC

36MG/IMPLANT

N020088 001 Dec 10, 1990

TABLET; ORAL

LEVONORGESTREL

ALVOGEN

1.5MG

A202246 001 Jun 05, 2015

FDN CONSUMER

0.75MG **

A078665 001 Aug 28, 2009

1.5MG

A200670 001 Jul 12, 2012

L PERRIGO CO

0.75MG

A090740 001 Dec 30, 2010

LOTUS PHARM CO LTD

0.75MG

A202684 001 Sep 02, 2016

LUPIN LTD

0.75MG

A091328 001 Jan 23, 2013

WATSON LABS

0.75MG

A078666 001 Jun 24, 2009

XIROMED

0.75MG

A202740 001 Sep 02, 2016

PLAN B

+ FDN CONSUMER

0.75MG **

N021045 001 Jul 28, 1999

+

0.75MG **

N021045 002 Aug 24, 2006

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

LEVOPROPOXYPHENE NAPSYLATE ANHYDROUS

CAPSULE; ORAL

NOVRAD

LILLY	EQ 50MG BASE	N012928	006
	EQ 100MG BASE	N012928	004

SUSPENSION; ORAL

NOVRAD

LILLY	EQ 50MG BASE/5ML	N012928	002
-------	------------------	---------	-----

LEVORPHANOL TARTRATE

INJECTABLE; INJECTION

LEVO-DROMORAN

VALEANT PHARM INTL	2MG/ML	N008719	001	Dec 19, 1991
--------------------	--------	---------	-----	--------------

TABLET; ORAL

LEVO-DROMORAN

+ VALEANT PHARM INTL	2MG **	N008720	001	Dec 19, 1991
----------------------	--------	---------	-----	--------------

LEVORPHANOL TARTRATE

HIKMA	1MG	A074278	002	Jun 18, 2018
	3MG	A074278	003	Jun 18, 2018

LEVOTHYROXINE SODIUM

CAPSULE; ORAL

LEVOTHYROXINE SODIUM

TEVA PHARMS USA INC	0.075MG	A211369	001	Oct 28, 2020
	0.088MG	A213256	001	Jan 06, 2021
	0.1MG	A213256	002	Jan 06, 2021
	0.112MG	A211369	003	Apr 16, 2021
	0.125MG	A213256	003	Jan 06, 2021
	0.137MG	A211369	005	May 02, 2023
	0.15MG	A211369	002	Oct 28, 2020
	0.175MG	A211369	006	May 02, 2023
	0.2MG	A211369	004	Nov 09, 2022

POWDER; INTRAVENOUS

LEVOTHYROXINE SODIUM

DR REDDYS	100MCG/VIAL	A208837	001	Mar 27, 2020
ENDO OPERATIONS	200MCG/VIAL	A205366	001	Dec 07, 2015

TABLET; ORAL

EUTHYROX

PROVELL	0.3MG	N021292	012	May 31, 2002
---------	-------	---------	-----	--------------

LEVOTHYROXINE SODIUM

AMNEAL	0.025MG	A210831	001	Feb 19, 2019
	0.05MG	A210831	002	Feb 19, 2019
	0.075MG	A210831	003	Feb 19, 2019
	0.088MG	A210831	004	Feb 19, 2019
	0.1MG	A210831	005	Feb 19, 2019
	0.112MG	A210831	006	Feb 19, 2019
	0.125MG	A210831	007	Feb 19, 2019
	0.137MG	A210831	008	Feb 19, 2019
	0.15MG	A210831	009	Feb 19, 2019
	0.175MG	A210831	010	Feb 19, 2019
	0.2MG	A210831	011	Feb 19, 2019
	0.3MG	A210831	012	Feb 19, 2019
MERCK KGAA	0.025MG	A076752	001	Jun 16, 2005
	0.05MG	A076752	002	Jun 16, 2005
	0.075MG	A076752	003	Jun 16, 2005
	0.088MG	A076752	004	Jun 16, 2005
	0.1MG	A076752	005	Jun 16, 2005
	0.112MG	A076752	006	Jun 16, 2005
	0.125MG	A076752	007	Jun 16, 2005
	0.15MG	A076752	008	Jun 16, 2005
	0.175MG	A076752	009	Jun 16, 2005
	0.2MG	A076752	010	Jun 16, 2005
	0.3MG	A076752	011	Jun 16, 2005

LEVOXYL

+ KING PHARMS	0.3MG **	N021301	012	May 25, 2001
---------------	----------	---------	-----	--------------

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

LIDOCAINE

AEROSOL;ORAL

XYLOCAINE

ASTRAZENECA 10% N014394 001

FILM, EXTENDED RELEASE;BUCCAL

DENTIPATCH

NOVEN 23MG/PATCH N020575 001 May 21, 1996

OINTMENT;TOPICAL

ALPHACAINE

CARLISLE 5% A084944 001

5% A084946 001

5% A084947 001

LIDOCAINE

BELMORA LLC 5% A080210 001

COSETTE 5% A211019 001 Dec 12, 2018

ENCUBE 5% A205318 001 Feb 01, 2016

GENEYORK PHARMS 5% A212486 001 Oct 17, 2019

RISING 5% A208604 001 Sep 20, 2017

TEVA PHARMS USA 5% A210256 001 Jan 16, 2018

VITRUVIAS THERAP 5% A208822 001 Sep 25, 2017

XYLOCAINE

+ ASTRAZENECA 5% ** N008048 001

PATCH;TOPICAL

DENTIPATCH

NOVEN 46.1MG/PATCH N020575 002 May 21, 1996

LIDOCAINE

NOVEN PHARMS INC 5% A203265 001 Dec 01, 2020

SOLUTION;TOPICAL

XYLOCAINE

ASTRAZENECA 5% N014127 001

SUPPOSITORY;RECTAL

XYLOCAINE

ASTRAZENECA 100MG N013077 001

LIDOCAINE HYDROCHLORIDE

INJECTABLE;INJECTION

ALPHACAINE HYDROCHLORIDE

CARLISLE 2% A084721 001

LIDOCAINE HYDROCHLORIDE

ABBOTT 10% A087980 001 Feb 02, 1983

20% A089362 001 May 25, 1988

ABRAXIS PHARM 1% A080420 001

1% A086761 001

1.5% A080420 005

2% A080420 002

2% A080420 004

2% A086761 002

2% N017508 001

4% N017508 002

20% N017508 004

AM REGENT 1% A080850 001

1% A091564 001 Aug 14, 2015

BEL MAR 1% A080710 001

2% A080760 001

BELMORA LLC 2% A080504 001

DELL LABS 1% A083387 001

2% A083388 001

ELKINS SINN 0.5% A085131 001

4% A084626 001

EPIC PHARMA LLC 1% A085037 001

2% A085037 002

GD SEARLE LLC 1% A083135 001

2% A083135 002

HOSPIRA 1% A040013 001 Jun 23, 1995

1.5% A088330 001 May 17, 1984

2% A088331 001 May 17, 1984

20% A083158 003

INTL MEDICATION 1% N017701 002

2% N017701 001

1GM/VIAL N018543 001

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

LIDOCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

LIDOCAINE HYDROCHLORIDE	2GM/VIAL	N018543	002	
LUITPOLD	2%	A083198	001	
LYPHOMED	1%	A080390	001	
	2%	A080390	002	
MILES	1%	A080414	001	
	2%	A080414	002	
RISING	0.5%	A091056	001	Dec 08, 2010
	0.5%	A091058	001	Sep 30, 2010
	1%	A091056	002	Dec 08, 2010
	1%	A091058	002	Sep 30, 2010
	2%	A202242	001	Apr 11, 2014
WATSON LABS	1%	A080377	001	
	1%	A083627	001	
	2%	A080377	002	
	2%	A083627	002	
WYETH AYERST	1%	A083083	001	
	2%	A083083	002	
LIDOCAINE HYDROCHLORIDE 0.1% AND DEXTROSE 5% IN PLASTIC CONTAINER				
BAXTER HLTHCARE	100MG/100ML	N018461	001	
LIDOCAINE HYDROCHLORIDE 0.2% AND DEXTROSE 5% IN PLASTIC CONTAINER				
B BRAUN	200MG/100ML	N018967	001	Mar 30, 1984
LIDOCAINE HYDROCHLORIDE 0.2% IN DEXTROSE 5%				
HOSPIRA	200MG/100ML	A083158	005	
LIDOCAINE HYDROCHLORIDE 0.2% IN DEXTROSE 5% IN PLASTIC CONTAINER				
ABBOTT	200MG/100ML	N018954	001	Jul 09, 1985
HOSPIRA	200MG/100ML	N018388	001	
LIDOCAINE HYDROCHLORIDE 0.4% AND DEXTROSE 5% IN PLASTIC CONTAINER				
B BRAUN	400MG/100ML	N018967	002	Mar 30, 1984
LIDOCAINE HYDROCHLORIDE 0.4% IN DEXTROSE 5%				
HOSPIRA	400MG/100ML	A083158	006	
LIDOCAINE HYDROCHLORIDE 0.4% IN DEXTROSE 5% IN PLASTIC CONTAINER				
HOSPIRA	400MG/100ML	N018388	002	
LIDOCAINE HYDROCHLORIDE 0.8% AND DEXTROSE 5% IN PLASTIC CONTAINER				
B BRAUN	800MG/100ML	N018967	003	Mar 30, 1984
LIDOCAINE HYDROCHLORIDE 0.8% IN DEXTROSE 5% IN PLASTIC CONTAINER				
HOSPIRA	800MG/100ML	N018388	003	Nov 05, 1982
LIDOCAINE HYDROCHLORIDE IN PLASTIC CONTAINER				
HOSPIRA	1.5%	A088326	001	Jul 31, 1984
	10%	A088367	001	Jul 31, 1984
	20%	A088368	001	Jul 31, 1984
LIDOCAINE HYDROCHLORIDE PRESERVATIVE FREE				
EXTROVIS	2%	A090665	001	Sep 27, 2010
INTL MEDICATION	4%	N017702	002	
	20%	N017702	001	
LIDOCATON				
PHARMATON	2%	A084727	001	Aug 17, 1983
LIDOPEN				
MERIDIAN MEDCL TECHN	10%	N017549	001	
XYLOCAINE				
ASTRAZENECA	1%	N010418	005	
	1.5%	N010418	009	
	2%	N010418	007	
XYLOCAINE 4% PRESERVATIVE FREE				
+ FRESENIUS KABI USA	4%	N010417	001	
XYLOCAINE DENTAL				
DENTSPLY PHARM	2%	N021380	001	
XYLOCAINE PRESERVATIVE FREE				
+ FRESENIUS KABI USA	1% **	N016801	005	Jan 19, 1988
+ FRESENIUS KABI USA	2% **	N016801	001	
+ FRESENIUS KABI USA	4% **	N016801	002	
+ FRESENIUS KABI USA	10% **	N016801	003	
+ FRESENIUS KABI USA	20% **	N016801	004	
INJECTABLE; SPINAL				
XYLOCAINE 1.5% W/ DEXTROSE 7.5%				
FRESENIUS KABI USA	1.5%	N016297	001	

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

LIDOCAINE HYDROCHLORIDE

INJECTABLE;SPINAL

XYLOCAINE 5% W/ GLUCOSE 7.5%

ASTRAZENECA 5%

N010496 002 Jul 07, 1982

JELLY;TOPICAL

ANESTACON

BIONPHARMA 2%

A080429 001

LIDOCAINE HYDROCHLORIDE

COSETTE 2%

A081318 001 Apr 29, 1993

WATSON LABS INC 2%

A040837 001 Mar 23, 2011

XYLOCAINE

+ EPIC PHARMA LLC 2% **

N008816 001

SOLUTION;ORAL

LIDOCAINE HYDROCHLORIDE

HIKMA 2%

A040014 001 Jul 10, 1995

LIDOCAINE HYDROCHLORIDE VISCOUS

ACTAVIS MID ATLANTIC 2%

A086578 001

INTL MEDICATION 2%

A086389 001 Feb 02, 1982

XYLOCAINE VISCOUS

+ FRESENIUS KABI USA 2% **

N009470 001

SOLUTION;TOPICAL

LARYNGOTRACHEAL ANESTHESIA KIT

KENDALL IL 4%

A087931 001 Jun 10, 1983

LIDOCAINE HYDROCHLORIDE

PACO 4%

A089688 001 Jun 30, 1989

LTA II KIT

HOSPIRA 4%

A080409 001

4%

A088542 001 Jul 31, 1984

PEDIATRIC LTA KIT

ABBOTT 2%

A088572 001 Jul 31, 1984

HOSPIRA 2%

A085995 001

XYLOCAINE 4% PRESERVATIVE FREE

+ FRESENIUS KABI USA 4% **

N010417 002

LIDOCAINE HYDROCHLORIDE; OXYTETRACYCLINE

INJECTABLE;INJECTION

TERRAMYCIN

PFIZER 2%;50MG/ML

A060567 001

2%;125MG/ML

A060567 002

LIDOCAINE; PRILOCAINE

CREAM;TOPICAL

EMLA

+ TEVA BRANDED PHARM 2.5%;2.5% **

N019941 001 Dec 30, 1992

LIDOCAINE AND PRILOCAINE

HIKMA 2.5%;2.5%

A076290 001 Sep 25, 2003

RHODES PHARMS 2.5%;2.5%

A213253 001 Sep 21, 2020

DISC;TOPICAL

EMLA

ASTRAZENECA 2.5%;2.5%

N020962 001 Feb 04, 1998

LIDOCAINE; TETRACAINE

CREAM;TOPICAL

PLIAGLIS

+ CRESCITA THERAP 7%;7%

N021717 001 Jun 29, 2006

PATCH;TOPICAL

SYNERA

+ GALEN SPECIALTY 70MG;70MG

N021623 001 Jun 23, 2005

LIFITEGRAST

SOLUTION/DROPS;OPHTHALMIC

LIFITEGRAST

EUGIA PHARMA 5%

A215063 001 Nov 07, 2023

INGENUS PHARMS LLC 5%

A215058 001 Oct 03, 2024

MICRO LABS 5%

A215081 001 Aug 04, 2023

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

LINACLOTIDE

CAPSULE; ORAL

LINACLOTIDE

AUROBINDO PHARMA	145MCG	A209611 001	Feb 07, 2023
	290MCG	A209611 002	Feb 07, 2023
MYLAN	145MCG	A209564 001	Feb 09, 2021
	290MCG	A209564 002	Feb 09, 2021

LINAGLIPTIN

TABLET; ORAL

LINAGLIPTIN

ZYDUS PHARMS	5MG	A208448 001	Mar 30, 2023
--------------	-----	-------------	--------------

LINAGLIPTIN; METFORMIN HYDROCHLORIDE

TABLET; ORAL

LINAGLIPTIN AND METFORMIN HYDROCHLORIDE

ZYDUS PHARMS	2.5MG; 500MG	A208449 001	Mar 30, 2023
	2.5MG; 850MG	A208449 002	Mar 30, 2023
	2.5MG; 1GM	A208449 003	Mar 30, 2023

LINCOMYCIN HYDROCHLORIDE

CAPSULE; ORAL

LINCOCIN

PHARMACIA AND UPJOHN	EQ 250MG BASE	N050316 001	
	EQ 500MG BASE	N050316 002	

INJECTABLE; INJECTION

LINCOMYCIN HYDROCHLORIDE

SLATE RUN PHARMA	EQ 300MG BASE/ML	A216662 001	Dec 21, 2022
SLAYBACK	EQ 300MG BASE/ML	A063180 001	Apr 16, 1991

LINDANE

CREAM; TOPICAL

KWELL

REED AND CARNRICK	1%	A084218 001	
	1%	N006309 001	

LOTION; TOPICAL

GAMENE

SOLA BARNES HIND	1%	A084989 001	
------------------	----	-------------	--

KWELL

REED AND CARNRICK	1%	A084218 002	
	1%	N006309 003	

LINDANE

OLTA PHARMS	1%	A087313 001	
-------------	----	-------------	--

WOCKHARDT BIO AG	1%	A088190 001	Aug 16, 1984
------------------	----	-------------	--------------

SCABENE

STIEFEL	1%	A086769 001	
---------	----	-------------	--

SHAMPOO; TOPICAL

GAMENE

SOLA BARNES HIND	1%	A084988 001	
------------------	----	-------------	--

KWELL

REED AND CARNRICK	1%	A084219 001	
	1%	N010718 001	

LINDANE

OLTA PHARMS	1%	A087266 001	
-------------	----	-------------	--

WOCKHARDT BIO AG	1%	A088191 001	Sep 18, 1984
------------------	----	-------------	--------------

SCABENE

STIEFEL	1%	A087940 001	Apr 08, 1983
---------	----	-------------	--------------

LINEZOLID

SOLUTION; INTRAVENOUS

LINEZOLID

HOSPIRA	600MG/300ML (2MG/ML)	A205442 001	Jul 07, 2015
---------	----------------------	-------------	--------------

TEVA PHARMS	600MG/300ML (2MG/ML)	A200222 001	Jun 27, 2012
-------------	----------------------	-------------	--------------

ZYVOX

+ PFIZER	400MG/200ML (2MG/ML) **	N021131 002	Apr 18, 2000
----------	-------------------------	-------------	--------------

TABLET; ORAL

LINEZOLID

AMNEAL PHARMS	600MG	A204536 001	Dec 21, 2015
---------------	-------	-------------	--------------

GATE PHARMS	600MG	A091210 001	Feb 05, 2016
-------------	-------	-------------	--------------

TEVA PHARMS USA	600MG	A078061 001	May 18, 2015
-----------------	-------	-------------	--------------

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

LINEZOLID

TABLET; ORAL

ZYVOX

+ PFIZER

400MG **

N021130 001 Apr 18, 2000

LIOTHYRONINE SODIUM

INJECTABLE; INJECTION

TRIOSTAT

+ ENDO OPERATIONS

EQ 0.01MG BASE/ML **

N020105 001 Dec 31, 1991

TABLET; ORAL

LIOTHYRONINE SODIUM

NORVIUM BIOSCIENCE

EQ 0.005MG BASE

A090326 001 Jul 14, 2009

EQ 0.025MG BASE

A090326 002 Jul 14, 2009

EQ 0.05MG BASE

A090326 003 Jul 14, 2009

WATSON LABS

EQ 0.025MG BASE

A085755 001 Jan 25, 1982

EQ 0.05MG BASE

A085753 001 Feb 03, 1982

LIOTRIX (T4;T3)

TABLET; ORAL

EUTHROID-0.5

PARKE DAVIS

0.03MG;0.0075MG

N016680 001

EUTHROID-1

PARKE DAVIS

0.06MG;0.015MG

N016680 002

EUTHROID-2

PARKE DAVIS

0.12MG;0.03MG

N016680 003

EUTHROID-3

PARKE DAVIS

0.18MG;0.045MG

N016680 004

THYROLAR-0.25

+ ALLERGAN

0.0125MG;0.0031MG

N016807 001

THYROLAR-0.5

+ ALLERGAN

0.025MG;0.0063MG

N016807 005

THYROLAR-1

+ ALLERGAN

0.05MG;0.0125MG

N016807 004

THYROLAR-2

+ ALLERGAN

0.1MG;0.025MG

N016807 002

THYROLAR-3

+ ALLERGAN

0.15MG;0.0375MG

N016807 003

THYROLAR-5

ALLERGAN

0.25MG;0.0625MG

N016807 006

LISDEXAMFETAMINE DIMESYLATE

CAPSULE; ORAL

LISDEXAMFETAMINE DIMESYLATE

SANDOZ

20MG

A202836 001 Jul 26, 2024

30MG

A202836 002 Jul 26, 2024

40MG

A202836 003 Jul 26, 2024

50MG

A202836 004 Jul 26, 2024

60MG

A202836 005 Jul 26, 2024

70MG

A202836 006 Jul 26, 2024

LISINAPRIL

TABLET; ORAL

LISINAPRIL

ACCORD HLTHCARE

2.5MG

A202554 001 Jul 30, 2013

5MG

A202554 002 Jul 30, 2013

10MG

A202554 003 Jul 30, 2013

20MG

A202554 004 Jul 30, 2013

30MG

A202554 005 Jul 30, 2013

40MG

A202554 006 Jul 30, 2013

HERITAGE PHARMA

2.5MG

A075752 001 Jul 01, 2002

5MG

A075752 002 Jul 01, 2002

10MG

A075752 003 Jul 01, 2002

20MG

A075752 004 Jul 01, 2002

30MG

A075752 005 Jul 01, 2002

40MG

A075752 006 Jul 01, 2002

HIKMA INTL PHARMS

2.5MG

A076063 001 Jul 01, 2002

5MG

A076063 002 Jul 01, 2002

10MG

A076063 003 Jul 01, 2002

20MG

A076063 004 Jul 01, 2002

30MG

A076063 006 Jun 27, 2003

40MG

A076063 005 Jul 01, 2002

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

LISINOPRIL

TABLET;ORAL

LISINOPRIL

RISING	2.5MG	A075999 001	Jul 01, 2002
	5MG	A075999 002	Jul 01, 2002
	10MG	A075999 003	Jul 01, 2002
	20MG	A075999 004	Jul 01, 2002
	30MG	A075999 005	Jul 01, 2002
	40MG	A075999 006	Jul 01, 2002
STRIDES PHARMA	2.5MG	A076071 001	Jul 01, 2002
	5MG	A076071 002	Jul 01, 2002
	10MG	A076071 003	Jul 01, 2002
	20MG	A076071 004	Jul 01, 2002
	30MG	A076071 005	Jul 01, 2002
	40MG	A076071 006	Jul 01, 2002
TEVA	2.5MG	A075783 001	Jul 01, 2002
	5MG	A075783 002	Jul 01, 2002
	10MG	A075783 003	Jul 01, 2002
	20MG	A075783 004	Jul 01, 2002
	30MG	A075783 005	Jul 01, 2002
	40MG	A075783 006	Jul 01, 2002
PRINIVIL			
MERCK	2.5MG	N019558 006	Jan 28, 1994
	5MG	N019558 001	Dec 29, 1987
	10MG	N019558 002	Dec 29, 1987
	20MG	N019558 003	Dec 29, 1987
	40MG	N019558 004	Oct 25, 1988

LITHIUM CARBONATE

CAPSULE;ORAL

ESKALITH

NOVEN THERAP	300MG	N016860 001	
--------------	-------	-------------	--

LITHIUM CARBONATE

ABLE	150MG	A076823 001	Jun 29, 2004
	300MG	A076121 001	Sep 27, 2001
	300MG	A076823 002	Jun 29, 2004
	600MG	A076823 003	Jun 29, 2004
APOTEX INC	300MG	A076795 001	Nov 22, 2004
NORVIUM BIOSCIENCE	150MG	A076243 002	Feb 24, 2003
	300MG	A076243 001	Jun 27, 2002
	600MG	A078763 001	Apr 15, 2008
USL PHARMA	300MG	A072542 001	Feb 01, 1989
WATSON LABS	300MG	A070407 001	Mar 19, 1987

LITHONATE

SOLVAY	300MG	N016782 001	
--------	-------	-------------	--

TABLET;ORAL

ESKALITH

JDS PHARMS	300MG	N017971 001	
------------	-------	-------------	--

LITHANE

BAYER PHARMS	300MG	N018833 001	Jul 18, 1985
--------------	-------	-------------	--------------

LITHIUM CARBONATE

HIKMA INTL PHARMS	300MG	A078715 001	Dec 28, 2010
PFIZER	300MG	N016834 001	

LITHOTABS

SOLVAY	300MG	N016980 001	
--------	-------	-------------	--

TABLET, EXTENDED RELEASE;ORAL

ESKALITH CR

JDS PHARMS	450MG **	N018152 001	Mar 29, 1982
------------	----------	-------------	--------------

LITHIUM CARBONATE

ABLE	300MG	A076382 001	Apr 21, 2003
ALEMBIC	300MG	A204445 001	Jun 10, 2015
HERITAGE PHARMA	300MG	A076170 001	Jun 10, 2002
	450MG	A076366 001	Aug 21, 2003
HIKMA INTL PHARMS	450MG	A076490 001	Jun 17, 2003

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

LITHIUM CITRATE

SYRUP;ORAL

LITHIUM CITRATE

+	HIKMA	EQ 300MG CARBONATE/5ML **	N018421	001	
	PAI HOLDINGS PHARM	EQ 300MG CARBONATE/5ML	A070755	001	May 21, 1986

LITHONATE

	SOLVAY	EQ 300MG CARBONATE/5ML	N017672	001	
--	--------	------------------------	---------	-----	--

LOMEFLOXACIN HYDROCHLORIDE

TABLET;ORAL

MAXAQUIN

	PHARMACIA	EQ 400MG BASE	N020013	001	Feb 21, 1992
--	-----------	---------------	---------	-----	--------------

LOMITAPIDE MESYLATE

CAPSULE;ORAL

JUXTAPID

+	CHIESI	EQ 40MG BASE	N203858	005	Apr 23, 2015
+		EQ 60MG BASE	N203858	006	Apr 23, 2015

LOMUSTINE

CAPSULE;ORAL

GLEOSTINE

+	LATINA PHARMA	5MG	N017588	004	Dec 19, 2014
---	---------------	-----	---------	-----	--------------

LOPERAMIDE HYDROCHLORIDE

CAPSULE;ORAL

IMODIUM

	J AND J CONSUMER INC	2MG **	N017690	001	
+		2MG **	N017694	001	

LOPERAMIDE HYDROCHLORIDE

	CHARTWELL RX	2MG	A072993	001	Aug 28, 1992
	ROXANE	2MG	A073080	001	Nov 27, 1991
	TEVA	2MG	A073122	001	Aug 30, 1991

SOLUTION;ORAL

IMODIUM

	JANSSEN PHARMS	1MG/5ML	N019037	001	Jul 31, 1984
--	----------------	---------	---------	-----	--------------

LOPERAMIDE HYDROCHLORIDE

	ALLIED	1MG/5ML	A073079	001	Apr 30, 1992
	ALPHARMA US PHARMS	1MG/5ML	A073187	001	Sep 15, 1992
	DURAMED PHARMS BARR	1MG/5ML	A074991	001	Dec 29, 1997
	PERRIGO	1MG/5ML	A073243	001	Jan 21, 1992
	SCIEGEN PHARMS INC	1MG/5ML	A074352	001	Nov 17, 1995
	TEVA	1MG/5ML	A073478	001	Jun 23, 1995
	WATSON LABS	1MG/5ML	A073062	001	May 28, 1993

TABLET;ORAL

LOPERAMIDE HYDROCHLORIDE

	ABLE	2MG	A073528	001	Nov 30, 1993
	CONTRACT PHARMACAL	2MG	A073254	001	Jul 30, 1993
	PERRIGO	2MG	A074194	001	Oct 30, 1992

TABLET, CHEWABLE;ORAL

IMODIUM A-D EZ CHEWS

+	J AND J CONSUMER INC	2MG	N020448	001	Jul 24, 1997
---	----------------------	-----	---------	-----	--------------

LOPERAMIDE HYDROCHLORIDE; SIMETHICONE

TABLET;ORAL

LOPERAMIDE HYDROCHLORIDE AND SIMETHICONE

	GRANULES	2MG;125MG	A215981	001	Aug 29, 2022
	SUN PHARM INDS LTD	2MG;125MG	A077500	001	Sep 06, 2006

TABLET, CHEWABLE;ORAL

IMODIUM MULTI-SYMPOM RELIEF

+	J AND J CONSUMER INC	2MG;125MG	N020606	001	Jun 26, 1996
---	----------------------	-----------	---------	-----	--------------

LOPERAMIDE HYDROCHLORIDE AND SIMETHICONE

	PERRIGO	2MG;125MG	A076029	001	Aug 30, 2002
--	---------	-----------	---------	-----	--------------

LOPINAVIR; RITONAVIR

CAPSULE;ORAL

KALETRA

	ABBVIE	133.3MG;33.3MG	N021226	001	Sep 15, 2000
--	--------	----------------	---------	-----	--------------

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

LORACARBEF

CAPSULE; ORAL

LORABID

KING PHARMS

200MG

N050668 001 Dec 31, 1991

400MG

N050668 002 Apr 05, 1996

FOR SUSPENSION; ORAL

LORABID

KING PHARMS

100MG/5ML

N050667 001 Dec 31, 1991

200MG/5ML

N050667 002 Dec 31, 1991

LORATADINE

CAPSULE; ORAL

LORATADINE

AUROBINDO PHARMA

10MG

A211900 001 Mar 24, 2023

STRIDES SOFTGELS

10MG

A211926 001 Jan 15, 2020

SYRUP; ORAL

CLARITIN HIVES RELIEF

+ BAYER HEALTHCARE LLC

1MG/ML **

N020641 003 Nov 19, 2003

LORATADINE

PHARM ASSOC

1MG/ML

A075565 001 Oct 05, 2004

RANBAXY LABS LTD

1MG/ML

A076529 001 Aug 20, 2004

TEVA

1MG/ML

A075505 001 Nov 07, 2003

TABLET; ORAL

LORATADINE

NORVIUM BIOSCIENCE

10MG

A075790 001 Nov 07, 2008

10MG

A078447 001 Aug 12, 2011

PERRIGO

10MG

N021512 001 Jun 24, 2004

TABLET, ORALLY DISINTEGRATING; ORAL

LORATADINE

ACTAVIS LABS FL INC

10MG

A075990 001 Nov 03, 2003

GLAXOSMITHKLINE

10MG

A075822 001 Feb 10, 2003

LORATADINE; PSEUDOEPHEDRINE SULFATE

TABLET, EXTENDED RELEASE; ORAL

LORATADINE AND PSEUDOEPHEDRINE SULFATE

HERITAGE PHARMA

5MG; 120MG

A076208 001 Jan 28, 2004

LORAZEPAM

CONCENTRATE; ORAL

LORAZEPAM

CHARTWELL MOLECULAR

2MG/ML

A079244 001 Apr 28, 2009

HIKMA

2MG/ML

A200169 001 Jan 30, 2012

INJECTABLE; INJECTION

LORAZEPAM

BEDFORD

2MG/ML

A077076 001 Jul 13, 2005

4MG/ML

A077076 002 Jul 13, 2005

DR REDDYS

1MG/0.5ML

A074551 003 Sep 12, 1996

2MG/ML

A074535 001 Sep 12, 1996

2MG/ML

A074551 001 Sep 12, 1996

4MG/ML

A074535 002 Sep 12, 1996

4MG/ML

A074551 002 Sep 12, 1996

ENDO OPERATIONS

2MG/ML

A074793 001 Mar 16, 2000

4MG/ML

A074793 002 Mar 16, 2000

EPIC PHARMA LLC

2MG/ML

A074974 001 Jul 23, 1998

HIKMA

2MG/ML

A074496 001 Sep 28, 1998

4MG/ML

A074496 002 Sep 28, 1998

HOSPIRA

2MG/ML

A074280 001 May 27, 1994

2MG/ML

A074300 001 Apr 12, 1994

4MG/ML

A074280 002 May 27, 1994

4MG/ML

A074300 003 Mar 19, 1997

RISING

2MG/ML

A200217 001 Apr 04, 2017

2MG/ML

A200542 001 Apr 28, 2017

4MG/ML

A200217 002 Apr 04, 2017

4MG/ML

A200542 002 Apr 28, 2017

WATSON LABS

2MG/ML

A074276 001 Apr 15, 1994

4MG/ML

A074276 002 Apr 15, 1994

LORAZEPAM PRESERVATIVE FREE

BEDFORD LABS

2MG/ML

A077074 001 Jul 13, 2005

4MG/ML

A077074 002 Jul 13, 2005

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

LORAZEPAM

SOLUTION; ORAL

LORAZEPAM

ROXANE

0.5MG/5ML

A074648 001 Mar 18, 1997

TABLET; ORAL

LORAZ

QUANTUM PHARMICS

0.5MG

A070200 001 Aug 09, 1985

1MG

A070201 001 Aug 09, 1985

2MG

A070202 001 Aug 09, 1985

LORAZEPAM

AM THERAP

0.5MG

A070727 001 Mar 07, 1986

1MG

A070728 001 Mar 07, 1986

2MG

A070729 001 Mar 07, 1986

AMNEAL PHARMS

0.5MG

A078826 001 Jun 23, 2010

1MG

A078826 002 Jun 23, 2010

2MG

A078826 003 Jun 23, 2010

ANDA REPOSITORY

0.5MG

A072555 002 Mar 29, 1991

1MG

A072555 003 Mar 29, 1991

2MG

A072555 001 Mar 29, 1991

CHARTWELL RX

0.5MG

A071591 002 Oct 13, 1987

1MG

A071591 003 Oct 13, 1987

2MG

A071591 001 Oct 13, 1987

HALSEY

0.5MG

A071434 001 Sep 01, 1987

1MG

A071435 001 Sep 01, 1987

2MG

A071436 001 Sep 01, 1987

MUTUAL PHARM

0.5MG

A070472 001 Dec 10, 1985

1MG

A070473 001 Dec 10, 1985

2MG

A070474 001 Dec 10, 1985

PAR PHARM

0.5MG

A070675 001 Dec 01, 1986

1MG

A070676 001 Dec 01, 1986

2MG

A070677 001 Dec 01, 1986

RISING

0.5MG

A077657 001 Mar 16, 2006

1MG

A077657 002 Mar 16, 2006

2MG

A077657 003 Mar 16, 2006

SANDOZ

0.5MG

A071193 001 Apr 15, 1988

1MG

A071194 001 Apr 15, 1988

2MG

A071195 001 Apr 15, 1988

SUPERPHARM

0.5MG

A071245 001 Feb 09, 1987

1MG

A071246 001 Feb 09, 1987

2MG

A071247 001 Feb 09, 1987

USL PHARMA

1MG

A070539 001 Dec 22, 1986

2MG

A070540 001 Dec 22, 1986

WARNER CHILCOTT

1MG

A071038 001 Jan 12, 1988

2MG

A071039 001 Jan 12, 1988

WATSON LABS

0.5MG

A071086 001 Mar 23, 1987

0.5MG

A071117 001 Jul 24, 1986

1MG

A071087 001 Mar 23, 1987

1MG

A071118 001 Jul 24, 1986

2MG

A071088 001 Mar 23, 1987

2MG

A071110 001 Jul 24, 1986

LOSARTAN POTASSIUM

TABLET; ORAL

LOSARTAN POTASSIUM

AIPING PHARM INC

25MG

A090544 001 Oct 06, 2010

50MG

A090544 002 Oct 06, 2010

100MG

A090544 003 Oct 06, 2010

APOTEX CORP

25MG

A090790 001 Oct 06, 2010

50MG

A090790 002 Oct 06, 2010

100MG

A090790 003 Oct 06, 2010

HIKMA

25MG

A077459 001 Oct 06, 2010

50MG

A077459 002 Oct 06, 2010

100MG

A077459 003 Oct 06, 2010

HISUN PHARM HANGZHOU

25MG

A204795 001 Apr 04, 2019

50MG

A204795 002 Apr 04, 2019

100MG

A204795 003 Apr 04, 2019

IPCA LABS LTD

25MG

A200290 001 Aug 30, 2013

50MG

A200290 002 Aug 30, 2013

100MG

A200290 003 Aug 30, 2013

MYLAN

25MG

A091590 001 Oct 06, 2010

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

LOSARTAN POTASSIUM

TABLET; ORAL

LOSARTAN POTASSIUM

	50MG	A091590 002	Oct 06, 2010
	100MG	A091590 003	Oct 06, 2010
TEVA	25MG	A076958 001	Apr 06, 2010
	50MG	A076958 002	Apr 06, 2010
	100MG	A076958 003	Apr 06, 2010
TORRENT PHARMS	25MG	A090467 001	Oct 06, 2010
	50MG	A090467 002	Oct 06, 2010
	100MG	A090467 003	Oct 06, 2010

LOTEPREDNOL ETABONATE

GEL; OPHTHALMIC

LOTEPREDNOL ETABONATE

HIKMA	0.5%	A213956 001	Nov 29, 2023
-------	------	-------------	--------------

SUSPENSION/DROPS; OPHTHALMIC

LOTEMAX

PHARMOS	0.5%	N020841 001	Mar 09, 1998
---------	------	-------------	--------------

LOVASTATIN

TABLET; ORAL

LOVASTATIN

AUROBINDO PHARMA USA	10MG	A075451 001	Dec 17, 2001
	10MG	A075935 001	Dec 17, 2001
	20MG	A075451 002	Dec 17, 2001
	20MG	A075935 002	Dec 17, 2001
	40MG	A075451 003	Dec 17, 2001
	40MG	A075935 003	Dec 17, 2001
SUN PHARM INDUSTRIES	10MG	A077520 001	Apr 14, 2006
	20MG	A077520 002	Apr 14, 2006
	40MG	A077520 003	Apr 14, 2006

MEVACOR

+ MERCK	10MG **	N019643 002	Mar 28, 1991
---------	---------	-------------	--------------

+	20MG **	N019643 003	Aug 31, 1987
---	---------	-------------	--------------

+	40MG **	N019643 004	Dec 14, 1988
---	---------	-------------	--------------

TABLET, EXTENDED RELEASE; ORAL

ALTOPREV

COVIS	10MG	N021316 001	Jun 26, 2002
-------	------	-------------	--------------

LOXAPINE HYDROCHLORIDE

CONCENTRATE; ORAL

LOXITANE C

TEVA BRANDED PHARM	EQ 25MG BASE/ML	N017658 001	
--------------------	-----------------	-------------	--

INJECTABLE; INJECTION

LOXITANE IM

ACTAVIS LABS UT INC	EQ 50MG BASE/ML	N018039 001	
---------------------	-----------------	-------------	--

LOXAPINE SUCCINATE

CAPSULE; ORAL

LOXAPINE SUCCINATE

RISING	EQ 5MG BASE	A076762 001	Nov 01, 2004
	EQ 10MG BASE	A076762 002	Nov 01, 2004
	EQ 25MG BASE	A076762 003	Nov 01, 2004
	EQ 50MG BASE	A076762 004	Nov 01, 2004

LOXITANE

+ TEVA BRANDED PHARM	EQ 5MG BASE **	N017525 001	
----------------------	----------------	-------------	--

+	EQ 10MG BASE **	N017525 002	
---	-----------------	-------------	--

+	EQ 25MG BASE **	N017525 003	
---	-----------------	-------------	--

+	EQ 50MG BASE **	N017525 004	
---	-----------------	-------------	--

TABLET; ORAL

LOXITANE

+ TEVA BRANDED PHARM	EQ 10MG BASE **	N017525 006	
----------------------	-----------------	-------------	--

+	EQ 25MG BASE **	N017525 007	
---	-----------------	-------------	--

+	EQ 50MG BASE **	N017525 008	
---	-----------------	-------------	--

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

LUBIPROSTONE

CAPSULE; ORAL

LUBIPROSTONE

ENDO OPERATIONS	8MCG	A201442 001	Jun 27, 2022
	24MCG	A201442 002	Jun 27, 2022

LUCINACTANT

SUSPENSION; INTRATRACHEAL

SURFAXIN

LEES PHARM HK	8.5ML	N021746 001	Mar 06, 2012
---------------	-------	-------------	--------------

LURASIDONE HYDROCHLORIDE

TABLET; ORAL

LURASIDONE HYDROCHLORIDE

TEVA PHARMS USA

	20MG	A208060 001	May 17, 2019
	40MG	A208060 002	May 17, 2019
	60MG	A208060 003	May 17, 2019
	80MG	A208060 004	May 17, 2019
	120MG	A208060 005	May 17, 2019
WATSON LABS TEVA	20MG	A208016 001	Feb 02, 2021
	40MG	A208016 002	Feb 02, 2021
	60MG	A208016 003	Feb 02, 2021
	80MG	A208016 004	Feb 02, 2021
	120MG	A208016 005	Feb 02, 2021

LYPRESSIN

SOLUTION; NASAL

DIAPID

NOVARTIS

0.185MG/ML

N016755 001

MACIMORELIN ACETATE

FOR SOLUTION; ORAL

MACRILEN

+ NOVO

EQ 60MG BASE/POUCH

N205598 001 Dec 20, 2017

MACITENTAN

TABLET; ORAL

MACITENTAN

APOTEX

10MG

A211195 001 Jan 09, 2024

AUROBINDO PHARMA LTD

10MG

A211198 001 Apr 18, 2023

TORRENT

10MG

A211107 001 Aug 05, 2024

ZYDUS LIFESCIENCES

10MG

A211224 001 Apr 06, 2021

MAFENIDE ACETATE

FOR SOLUTION; TOPICAL

MAFENIDE ACETATE

ENDO OPERATIONS

5%

A201511 001 Feb 12, 2013

NOVAST LABS

5%

A206716 001 Jul 31, 2017

SULFAMYLON

+ MYLAN INSTITUTIONAL

5%

N019832 003 Jun 05, 1998

MAGNESIUM ACETATE TETRAHYDRATE; POTASSIUM ACETATE; SODIUM CHLORIDE

INJECTABLE; INJECTION

PLASMA-LYTE 56 IN PLASTIC CONTAINER

BAXTER HLTHCARE

32MG/100ML; 128MG/100ML; 234MG/100ML

N019047 001 Jun 15, 1984

MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; POTASSIUM PHOSPHATE, MONOBASIC; SODIUM ACETATE; SODIUM CHLORIDE; SODIUM GLUCONATE; SODIUM PHOSPHATE, DIBASIC, HEPTAHYDRATE

INJECTABLE; INJECTION

ISOLYTE S PH 7.4 IN PLASTIC CONTAINER

B BRAUN

30MG/100ML; 37MG/100ML; 0.82MG/100ML; 370MG/100ML; 530MG/100ML; 500MG/100ML; 12MG/100ML

N019006 001 Apr 04, 1984

MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE; SODIUM GLUCONATE

INJECTABLE; INJECTION

ISOLYTE S IN PLASTIC CONTAINER

B BRAUN

30MG/100ML; 37MG/100ML; 370MG/100ML; 530MG/100ML; 500MG/100ML

N018252 001

SOLUTION; IRRIGATION

PHYSIOSOL IN PLASTIC CONTAINER

HOSPIRA INC

14MG/100ML; 37MG/100ML; 222MG/100ML; 526MG/100ML; 502MG/100ML

N018406 001

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE; SODIUM GLUCONATE

SOLUTION;IRRIGATION

PHYSIOSOL PH 7.4 IN PLASTIC CONTAINER

HOSPIRA INC	30MG/100ML;37MG/100ML;222MG/100ML;526MG /100ML;502MG/100ML	N018406 002	Jul 08, 1982
-------------	---	-------------	--------------

SYNOVALYTE IN PLASTIC CONTAINER

BAXTER HLTHCARE	30MG/100ML;37MG/100ML;368MG/100ML;526MG /100ML;502MG/100ML	N019326 001	Jan 25, 1985
-----------------	---	-------------	--------------

MAGNESIUM HYDROXIDE; OMEPRAZOLE; SODIUM BICARBONATE

TABLET;ORAL

MAGNESIUM HYDROXIDE AND OMEPRAZOLE AND SODIUM BICARBONATE

SANTARUS	343MG;20MG;750MG	N022456 001	Dec 04, 2009
	343MG;40MG;750MG	N022456 002	Dec 04, 2009

TABLET, CHEWABLE;ORAL

ZEGERID

SANTARUS	700MG;20MG;600MG	N021850 001	Mar 24, 2006
	700MG;40MG;600MG	N021850 002	Mar 24, 2006

MAGNESIUM SULFATE

SOLUTION;INTRAMUSCULAR, INTRAVENOUS

MAGNESIUM SULFATE

EXELA PHARMA	5GM/10ML (500MG/ML)	A206039 001	Dec 18, 2014
--------------	---------------------	-------------	--------------

MAGNESIUM SULFATE; POLYETHYLENE GLYCOL 3350; POTASSIUM CHLORIDE; POTASSIUM SULFATE; SODIUM BICARBONATE; SODIUM CHLORIDE; SODIUM SULFATE

SOLUTION;ORAL

SUCLEAR

+ BRAINTREE LABS	1.6GM/BOT, 3.13GM/BOT, 17.5GM/BOT, N/A, N/A , N/A, N/A; N/A, N/A, N/A, 210GM, 0.74GM, 2.86G M, 5.6GM **	N203595 001	Jan 18, 2013
------------------	---	-------------	--------------

MAGNESIUM SULFATE; POTASSIUM SULFATE; SODIUM SULFATE

POWDER;ORAL

COLPREP KIT

+ GATOR PHARMS	1.6GM/BOT;3.13GM/BOT;17.5GM/BOT	N204553 001	Dec 27, 2016
----------------	---------------------------------	-------------	--------------

MALATHION

LOTION;TOPICAL

MALATHION

NORVIUM BIOSCIENCE	0.5%	A078743 001	Mar 06, 2009
--------------------	------	-------------	--------------

OVIDE

+ TARO	0.5% **	N018613 001	Aug 02, 1982
--------	---------	-------------	--------------

MANGAFODIPIR TRISODIUM

INJECTABLE;INJECTION

TESLASCAN

IC TARGETS	37.9MG/ML	N020652 001	Nov 26, 1997
------------	-----------	-------------	--------------

MANGANESE CHLORIDE TETRAHYDRATE

FOR SOLUTION;ORAL

LUMENHANCE

BRACCO	3.49MG/GM	N020686 001	Dec 19, 1997
--------	-----------	-------------	--------------

MANGANESE SULFATE

INJECTABLE;INJECTION

MANGANESE SULFATE

+ ABRAXIS PHARM	EQ 0.1MG MANGANESE/ML **	N019228 001	May 05, 1987
-----------------	--------------------------	-------------	--------------

MANNITOL

INJECTABLE;INJECTION

MANNITOL 10%

B BRAUN	10GM/100ML	N016080 002	
---------	------------	-------------	--

HOSPIRA	10GM/100ML	N016269 002	
---------	------------	-------------	--

MILES	10GM/100ML	N016472 002	
-------	------------	-------------	--

MANNITOL 10% IN PLASTIC CONTAINER

ICU MEDICAL INC	10GM/100ML	N019603 002	Jan 08, 1987
-----------------	------------	-------------	--------------

MANNITOL 10% W/ DEXTROSE 5% IN DISTILLED WATER

B BRAUN	10GM/100ML	N016080 006	
---------	------------	-------------	--

MANNITOL 15%

B BRAUN	15GM/100ML	N016080 003	
---------	------------	-------------	--

HOSPIRA	15GM/100ML	N016269 003	
---------	------------	-------------	--

MILES	15GM/100ML	N016472 005	
-------	------------	-------------	--

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

MANNITOL

INJECTABLE; INJECTION

MANNITOL 15% IN PLASTIC CONTAINER			
ICU MEDICAL INC	15GM/100ML	N019603 003	Jan 08, 1990
MANNITOL 15% W/ DEXTROSE 5% IN SODIUM CHLORIDE 0.45%			
B BRAUN	15GM/100ML	N016080 005	
MANNITOL 20%			
B BRAUN	20GM/100ML	N014738 001	
	20GM/100ML	N016080 004	
HOSPIRA	20GM/100ML	N016269 004	
MILES	20GM/100ML	N016472 004	
MANNITOL 25%			
ABRAXIS PHARM	12.5GM/50ML	A086754 001	
HOSPIRA	12.5GM/50ML	N016269 005	
IGI LABS INC	12.5GM/50ML	A089239 001	May 06, 1987
	12.5GM/50ML	A089240 001	May 06, 1987
INTL MEDICATION	12.5GM/50ML	A083051 001	
LUITPOLD	12.5GM/50ML	A087409 001	Jan 21, 1982
MERCK	12.5GM/50ML	N005620 001	
WATSON LABS	12.5GM/50ML	A087460 001	Jun 27, 1983
MANNITOL 5%			
B BRAUN	5GM/100ML	N016080 001	
HOSPIRA	5GM/100ML	N016269 001	
MANNITOL 5% IN PLASTIC CONTAINER			
ICU MEDICAL INC	5GM/100ML	N019603 001	Jan 08, 1987
MANNITOL 5% W/ DEXTROSE 5% IN SODIUM CHLORIDE 0.12%			
B BRAUN	5GM/100ML	N016080 007	
SOLUTION; IRRIGATION			
RESECTISOL			
B BRAUN	5GM/100ML	N016704 002	
RESECTISOL IN PLASTIC CONTAINER			
B BRAUN	5GM/100ML	N016772 002	

MANNITOL; SORBITOL

SOLUTION; IRRIGATION

SORBITOL-MANNITOL			
HOSPIRA	540MG/100ML; 2.7GM/100ML	A080224 001	
SORBITOL-MANNITOL IN PLASTIC CONTAINER			
HOSPIRA	540MG/100ML; 2.7GM/100ML	N017636 001	

MAPROTILINE HYDROCHLORIDE

TABLET; ORAL

LUDIOMIL			
NOVARTIS	25MG	N017543 001	
	50MG	N017543 002	
	75MG	N017543 003	Sep 30, 1982
MAPROTILINE HYDROCHLORIDE			
AM THERAP	25MG	A072129 001	Jan 14, 1988
	50MG	A072130 001	Jan 14, 1988
	75MG	A072131 001	Jan 14, 1988
HERITAGE PHARMA	25MG	A072162 001	Jun 01, 1988
	50MG	A072163 001	Jun 01, 1988
RISING	25MG	A072285 002	Oct 03, 1988
	50MG	A072285 001	Oct 03, 1988
	75MG	A072285 003	Oct 03, 1988
WATSON LABS	25MG	A071943 001	Dec 30, 1987
	50MG	A071944 001	Dec 30, 1987
	75MG	A071945 001	Dec 30, 1987
	75MG	A072164 001	Jun 01, 1988

MARAVIROC

TABLET; ORAL

SELZENTRY			
+	VIIV HLTHCARE	25MG **	N022128 003 Nov 04, 2016
+		75MG **	N022128 004 Nov 04, 2016

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

MASOPROCOL

CREAM;TOPICAL

ACTINEX

UNIV AZ CANCER CTR 10%

N019940 001 Sep 04, 1992

MAZINDOL

TABLET;ORAL

MAZANOR

WYETH AYERST 1MG

N017980 002

2MG

N017980 001

SANOREX

+ NOVARTIS 1MG **

N017247 001

+ 2MG **

N017247 002

MEBENDAZOLE

TABLET, CHEWABLE;ORAL

VERMOX

+ JANSSEN PHARMS 100MG **

N017481 001

+ 500MG

N208398 001 Oct 19, 2016

MEBUTAMATE

TABLET;ORAL

DORMATE

MEDPOINTE PHARM HLC 600MG

N017374 001

MECAMYLAMINE HYDROCHLORIDE

TABLET;ORAL

INVERSINE

+ TARGACEPT 2.5MG **

N010251 001

MECHLORETHAMINE HYDROCHLORIDE

INJECTABLE; INJECTION

MUSTARGEN

+ RECORDATI RARE 10MG/VIAL

N006695 001

MECLIZINE HYDROCHLORIDE

TABLET;ORAL

MECLIZINE HYDROCHLORIDE

ABC HOLDING 12.5MG

A085253 001

25MG

A085252 001

ANABOLIC 25MG

A085891 001

ANI PHARMS 12.5MG

A084657 002

12.5MG

A085269 001

12.5MG

A088732 001 Dec 11, 1985

25MG

A084657 001

25MG

A085740 001

BUNDY 12.5MG

A084382 001

25MG

A084872 001

CHARTWELL RX 12.5MG

A205136 001 Feb 22, 2019

25MG

A205136 002 Feb 22, 2019

50MG

A205136 003 Feb 22, 2019

INVAGEN PHARMS 12.5MG

A200432 001 Feb 17, 2022

25MG

A200432 002 Feb 17, 2022

50MG

A200432 003 Feb 17, 2022

IVAX SUB TEVA PHARMS 12.5MG

A083784 001

KV PHARM 12.5MG

A085524 001

25MG

A085523 001

PLIVA 25MG

A088734 001 Dec 11, 1985

RISING 12.5MG

A040179 001 Jan 30, 1997

25MG

A040179 002 Jan 30, 1997

STRIDES PHARMA 50MG

A089674 001 Mar 31, 1988

SUPERPHARM 12.5MG

A089113 001 Aug 20, 1985

25MG

A089114 001 Aug 20, 1985

UDL 12.5MG

A088256 001 Jun 13, 1983

25MG

A088257 001 Jun 13, 1983

VANGARD 12.5MG

A087877 001 Apr 20, 1982

25MG

A087620 001 Jan 04, 1982

WATSON LABS 12.5MG

A085195 001

TABLET, CHEWABLE;ORAL

MECLIZINE HYDROCHLORIDE

INVAGEN PHARMS 25MG

A200791 001 Feb 17, 2022

IVAX SUB TEVA PHARMS 25MG

A084976 001

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

MECLIZINE HYDROCHLORIDE

TABLET, CHEWABLE;ORAL

MECLIZINE HYDROCHLORIDE

NEXGEN PHARMA INC 25MG

A086392 001

PLIVA 25MG

A088733 001 Dec 11, 1985

MECLOCYCLINE SULFOSALICYLATE

CREAM;TOPICAL

MECLAN

JOHNSON AND JOHNSON 1%

N050518 001

MECLOFENAMATE SODIUM

CAPSULE;ORAL

MECLODIUM

QUANTUM PHARMICS EQ 50MG BASE

A071380 001 Jul 14, 1987

EQ 100MG BASE

A071381 001 Jul 14, 1987

MECLOFENAMATE SODIUM

AM THERAP EQ 50MG BASE

A071362 001 Feb 10, 1987

EQ 100MG BASE

A071363 001 Feb 10, 1987

ANI PHARMS EQ 50MG BASE

A071469 002 Apr 15, 1987

EQ 100MG BASE

A071469 001 Apr 15, 1987

BARR EQ 50MG BASE

A072848 001 Mar 20, 1989

EQ 100MG BASE

A072809 001 Mar 20, 1989

CHARTWELL RX EQ 50MG BASE

A072262 001 Nov 29, 1988

EQ 100MG BASE

A072263 001 Nov 29, 1988

PAR PHARM EQ 50MG BASE

A072077 001 Mar 10, 1988

EQ 100MG BASE

A072078 001 Mar 10, 1988

USL PHARMA EQ 50MG BASE

A071007 001 Mar 25, 1988

EQ 100MG BASE

A071008 001 Mar 25, 1988

VITARINE EQ 50MG BASE

A071710 001 Jun 15, 1988

EQ 100MG BASE

A071684 001 Jun 15, 1988

WATSON LABS EQ 50MG BASE

A070400 001 Nov 25, 1986

EQ 50MG BASE

A071640 001 Aug 11, 1987

EQ 100MG BASE

A070401 001 Nov 25, 1986

EQ 100MG BASE

A071641 001 Aug 11, 1987

MECLOMEN

PARKE DAVIS EQ 50MG BASE

N018006 001

EQ 100MG BASE

N018006 002

MEDROXYPROGESTERONE ACETATE

INJECTABLE; INJECTION

DEPO-PROVERA

+ PFIZER 100MG/ML **

N012541 002

+ 400MG/ML

N012541 003

MEDROXYPROGESTERONE ACETATE

CIPLA 150MG/ML

A210335 001 Jan 25, 2019

HIKMA 150MG/ML

A214309 001 Jan 05, 2023

SANDOZ 150MG/ML

A078711 001 May 20, 2009

SUN PHARM 150MG/ML

A210760 001 May 01, 2019

150MG/ML

A210761 001 Apr 24, 2019

TEVA PHARMS USA 150MG/ML

A076552 001 Oct 27, 2004

TABLET;ORAL

AMEN

AMARIN PHARMS 10MG

A083242 001

CURRETAB

SOLVAY 10MG

A085686 001

CYCRIN

ESI 2.5MG

A081239 001 Oct 30, 1992

5MG

A081240 001 Oct 30, 1992

10MG

A089386 001 Sep 09, 1987

MEDROXYPROGESTERONE ACETATE

DURAMED PHARMS BARR 2.5MG

A040311 001 Dec 01, 1999

5MG

A040311 002 Dec 01, 1999

10MG

A040311 003 Dec 01, 1999

UPSHER SMITH LABS 10MG

A088484 001 Jul 26, 1984

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

MEDRYSONESUSPENSION;OPHTHALMIC
HMS

ALLERGAN 1% N016624 003

MEFENAMIC ACID

CAPSULE;ORAL

MEFENAMIC ACID

CHARTWELL RX 250MG A209209 001 Sep 18, 2020

NOSTRUM LABS INC 250MG A090359 001 Feb 05, 2013

MEFLOQUINE HYDROCHLORIDE

TABLET;ORAL

LARIAM

+ ROCHE 250MG ** N019591 001 May 02, 1989

MEFLOQUINE HYDROCHLORIDE

CHARTWELL RX 250MG A076175 001 Feb 20, 2002

HIKMA INTL PHARMS 250MG A077699 001 Apr 21, 2010

US ARMY WALTER REED 250MG ** N019578 001 May 02, 1989

MEGESTROL ACETATE

SUSPENSION;ORAL

MEGACE

+ BRISTOL MYERS SQUIBB 40MG/ML ** N020264 001 Sep 10, 1993

MEGESTROL ACETATE

CHARTWELL 125MG/ML A204688 001 Dec 01, 2017

HIKMA 40MG/ML A075997 001 Feb 15, 2002

40MG/ML A203960 001 Jun 09, 2017

TEVA PHARMS 40MG/ML A075681 001 May 05, 2003

XTTRIUM LABS INC 40MG/ML A076721 001 Nov 01, 2004

TABLET;ORAL

MEGACE

+ BRISTOL MYERS SQUIBB 20MG ** N016979 001

+ 40MG ** N016979 002

MEGESTROL ACETATE

HIKMA 20MG A074458 001 Sep 29, 1995

40MG A074458 002 Sep 29, 1995

TEVA 40MG A074745 001 Feb 27, 1998

USL PHARMA 20MG A070646 001 Oct 02, 1987

40MG A070647 001 Oct 02, 1987

MELOXICAM

CAPSULE;ORAL

VIVLODEX

+ ICEUTICA OPERATIONS 5MG ** N207233 001 Oct 22, 2015

+ 10MG ** N207233 002 Oct 22, 2015

SOLUTION;INTRAVENOUS

ANJESO

+ BAUDAX 30MG/ML (30MG/ML) ** N210583 001 Feb 20, 2020

TABLET;ORAL

MELOXICAM

ANDA REPOSITORY 7.5MG A077935 001 Jul 19, 2006

15MG A077935 002 Jul 19, 2006

CHARTWELL RX 7.5MG A077936 001 Jul 19, 2006

15MG A077936 002 Jul 19, 2006

CR DOUBLE CRANE 7.5MG A078039 001 Dec 14, 2006

15MG A078039 002 Dec 14, 2006

IMPAX LABS INC 7.5MG A077930 001 Jul 19, 2006

15MG A077930 002 Jul 19, 2006

MYLAN 7.5MG A077934 001 Jul 20, 2006

15MG A077934 002 Jul 20, 2006

NATCO PHARMA 7.5MG A077923 001 Jul 19, 2006

15MG A077923 002 Jul 19, 2006

ROXANE 7.5MG A077925 001 Jul 19, 2006

15MG A077925 002 Jul 19, 2006

SUN PHARM INDS INC 7.5MG A077937 001 Jul 19, 2006

15MG A077937 002 Jul 19, 2006

YABAO PHARM 7.5MG A077933 001 Jul 19, 2006

15MG A077933 002 Jul 19, 2006

MOBIC

+ BOEHRINGER INGELHEIM 7.5MG N020938 001 Apr 13, 2000

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

MELOXICAM

TABLET; ORAL

MOBIC

+ 15MG N020938 002 Aug 23, 2000

TABLET, ORALLY DISINTEGRATING; ORAL

QMIIZ ODT

+ TERSERA 7.5MG ** N211210 001 Oct 19, 2018

+ 15MG ** N211210 002 Oct 19, 2018

MELPHALAN

TABLET; ORAL

ALKERAN

+ APOTEX 2MG ** N014691 002

MELPHALAN

ALVOGEN 2MG A207809 001 Mar 22, 2017

MELPHALAN HYDROCHLORIDE

INJECTABLE; INJECTION

ALKERAN

+ APOTEX EQ 50MG BASE/VIAL ** N020207 001 Nov 18, 1992

MELPHALAN HYDROCHLORIDE

ENDO OPERATIONS EQ 50MG BASE/VIAL A204773 001 Aug 22, 2016

MYLAN INSTITUTIONAL EQ 50MG BASE/VIAL A090299 001 Oct 27, 2009

USWM EQ 50MG BASE/VIAL A207032 001 May 03, 2019

POWDER; INTRAVENOUS

MELPHALAN HYDROCHLORIDE

ACTAVIS LLC EQ 50MG BASE/VIAL A209323 001 Mar 06, 2020

MEMANTINE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

MEMANTINE HYDROCHLORIDE

ANI PHARMS 7MG A205365 001 Feb 28, 2020

14MG A205365 002 Feb 28, 2020

21MG A205365 003 Feb 28, 2020

28MG A205365 004 Feb 28, 2020

ENDO OPERATIONS 7MG A205784 001 Jun 09, 2017

14MG A205784 002 Jun 09, 2017

21MG A205784 003 Jun 09, 2017

28MG A205784 004 Jun 09, 2017

RISING 7MG A206032 001 Sep 28, 2016

14MG A206032 002 Sep 28, 2016

21MG A206032 003 Sep 28, 2016

28MG A206032 004 Sep 28, 2016

SUN PHARM 7MG A205905 001 Sep 28, 2016

14MG A205905 002 Sep 28, 2016

21MG A205905 003 Sep 28, 2016

28MG A205905 004 Sep 28, 2016

NAMENDA XR

+ ABBVIE 7MG ** N022525 001 Jun 21, 2010

+ 14MG N022525 002 Jun 21, 2010

+ 21MG N022525 003 Jun 21, 2010

+ 28MG N022525 004 Jun 21, 2010

SOLUTION; ORAL

MEMANTINE HYDROCHLORIDE

SCIEGEN PHARMS INC 2MG/ML A205446 001 Dec 07, 2015

NAMENDA

+ ALLERGAN 2MG/ML ** N021627 001 Apr 18, 2005

TABLET; ORAL

MEMANTINE HYDROCHLORIDE

ALLIED 5MG A090073 001 Sep 04, 2015

10MG A090073 002 Sep 04, 2015

CHARTWELL 5MG A090244 001 Jul 11, 2018

10MG A090244 002 Jul 11, 2018

HIKMA 5MG A208173 001 Feb 28, 2020

10MG A208173 002 Feb 28, 2020

JUBILANT GENERICS 5MG A091585 001 Oct 13, 2015

10MG A091585 002 Oct 13, 2015

LANNETT CO INC 5MG A207236 001 Nov 10, 2016

10MG A207236 002 Nov 10, 2016

ORBION PHARMS 5MG A090044 001 Mar 12, 2012

10MG A090044 002 Mar 12, 2012

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

MEMANTINE HYDROCHLORIDE

TABLET; ORAL

MEMANTINE HYDROCHLORIDE

RISING	5MG	A079225 001	Jan 30, 2015
	10MG	A079225 002	Jan 30, 2015
TEVA PHARMS	5MG	A090052 001	Oct 25, 2011
	10MG	A090052 002	Oct 25, 2011
TORRENT	5MG	A200155 001	Oct 13, 2015
	10MG	A200155 002	Oct 13, 2015
YILING	5MG	A212947 001	Apr 03, 2020
	10MG	A212947 002	Apr 03, 2020

MENADIOL SODIUM DIPHOSPHATE

INJECTABLE; INJECTION

KAPPADIONE

LILLY	10MG/ML	N005725 001	
SYNKAYVITE			
ROCHE	5MG/ML	N003718 004	
	10MG/ML	N003718 006	
	37.5MG/ML	N003718 008	

TABLET; ORAL

SYNKAYVITE

ROCHE	5MG	N003718 010	
-------	-----	-------------	--

MENADIONE

TABLET; ORAL

MENADIONE

LILLY	5MG	N002139 003	
-------	-----	-------------	--

MEPENZOLATE BROMIDE

SOLUTION; ORAL

CANTIL

SANOFI AVENTIS US	25MG/5ML	N010679 004	
-------------------	----------	-------------	--

TABLET; ORAL

CANTIL

+ SANOFI AVENTIS US	25MG	N010679 003	
---------------------	------	-------------	--

MEPERIDINE HYDROCHLORIDE

INJECTABLE; INJECTION

DEMEROL

QUAGEN	25MG/ML	N005010 007	
	50MG/ML	N005010 002	
	75MG/ML	N005010 009	
	100MG/ML	N005010 003	

MEPERIDINE HYDROCHLORIDE

ABBOTT

	25MG/ML	A080388 001	
	50MG/ML	A080385 001	
	50MG/ML	A080387 001	
	75MG/ML	A080389 001	
	100MG/ML	A080386 001	

BAXTER HLTHCARE

	25MG/ML	A088279 001	Jun 15, 1984
	50MG/ML	A088280 001	Jun 15, 1984
	75MG/ML	A088281 001	Jun 15, 1984
	100MG/ML	A088282 001	Jun 15, 1984

IGI LABS INC

	25MG/ML	A089781 001	Mar 31, 1989
	50MG/ML	A089782 001	Mar 31, 1989
	50MG/ML	A089783 001	Mar 31, 1989
	50MG/ML	A089784 001	Mar 31, 1989
	75MG/ML	A089785 001	Mar 31, 1989
	100MG/ML	A089786 001	Mar 31, 1989
	100MG/ML	A089787 001	Mar 31, 1989
	100MG/ML	A089788 001	Mar 31, 1989

INTL MEDICATION

PARKE DAVIS	10MG/ML	A086332 001	
	50MG/ML	A080364 002	
	75MG/ML	A080364 003	
	100MG/ML	A080364 001	

WATSON LABS

	50MG/ML	A073444 001	Mar 17, 1992
	100MG/ML	A073445 001	Mar 17, 1992

MEPERIDINE HYDROCHLORIDE PRESERVATIVE FREE

HOSPIRA

	10MG/ML	A040305 001	Mar 10, 1999
--	---------	-------------	--------------

+ ICU MEDICAL INC

	10MG/ML	A088432 001	Aug 16, 1984
--	---------	-------------	--------------

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

MEPERIDINE HYDROCHLORIDE

INJECTABLE; INJECTION

MEPERIDINE HYDROCHLORIDE PRESERVATIVE FREE

INTL MEDICATION	10MG/ML	A081309	001	Aug 30, 1993
SPECGX LLC	10MG/ML	A040163	001	May 12, 1997
WATSON LABS	10MG/ML	A073443	001	Mar 17, 1992

SYRUP; ORAL

DEMEROL

QUAGEN	50MG/5ML **	N005010	005	
--------	-------------	---------	-----	--

TABLET; ORAL

DEMEROL

+ QUAGEN	50MG **	N005010	001	
+	100MG	N005010	004	

MEPERIDINE HYDROCHLORIDE

BARR	50MG	A088639	001	Jul 02, 1984
	100MG	A088640	001	Sep 19, 1984
DURAMED PHARMS BARR	50MG	A040318	001	Oct 05, 1999
	100MG	A040318	002	Oct 05, 1999
GENUS	75MG	A040893	002	Jun 24, 2009
	100MG	A040893	003	Jun 24, 2009
	150MG	A040893	004	Jun 24, 2009
HIKMA	50MG	A040110	001	Mar 12, 1997
	100MG	A040110	002	Mar 12, 1997
SPECGX LLC	50MG	A040352	001	Jun 13, 2000
	100MG	A040352	002	Jun 13, 2000
STRIDES PHARMA	50MG	A040191	001	Dec 17, 1998
	100MG	A040191	002	Dec 17, 1998
SUN PHARM INDS INC	50MG	A040446	001	Aug 08, 2002
	100MG	A040446	002	Aug 08, 2002
SUN PHARM INDUSTRIES	50MG	A080448	001	
	100MG	A080448	002	
WATSON LABS	50MG	A040186	001	Jun 30, 1997
	100MG	A040186	002	Jun 30, 1997
WYETH AYERST	50MG	A080454	001	

MEPERIDINE HYDROCHLORIDE; PROMETHAZINE HYDROCHLORIDE

INJECTABLE; INJECTION

MEPERGAN

HIKMA	25MG/ML; 25MG/ML	N011730	001	
-------	------------------	---------	-----	--

MEPHENTERMINE SULFATE

INJECTABLE; INJECTION

WYAMINE SULFATE

BAXTER HLTHCARE CORP	EQ 15MG BASE/ML	N008248	002	
	EQ 30MG BASE/ML	N008248	001	

MEPHENYTOIN

TABLET; ORAL

MESANTOIN

+ NOVARTIS	100MG **	N006008	001	
------------	----------	---------	-----	--

MEPIVACAINE HYDROCHLORIDE

INJECTABLE; INJECTION

ARESTOCAINE HYDROCHLORIDE

SOLVAY	3%	A084777	002	Apr 18, 1982
--------	----	---------	-----	--------------

CARBOCAINE

+ EASTMAN KODAK	3% **	N012125	003	
-----------------	-------	---------	-----	--

ISOCAINE HYDROCHLORIDE

SEPTODONT	3%	A080925	001	
-----------	----	---------	-----	--

MEPIVACAINE HYDROCHLORIDE

BELMORA LLC	3%	A083559	001	
HOSPIRA INC	3%	A040806	001	Apr 28, 2008
INTL MEDICATION SYS	1%	A087509	001	Oct 05, 1982
WATSON LABS	1%	A088769	001	Nov 20, 1984
	2%	A088770	001	Nov 20, 1984

POLOCAINE

DENTSPLY PHARM	3%	A088653	001	Aug 21, 1984
----------------	----	---------	-----	--------------

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

MEPREDNISON

TABLET;ORAL

BETAPAR

SCHERING

4MG

N016053 002

MEPROBAMATE

CAPSULE;ORAL

EQUANIL

WYETH AYERST

400MG

N012455 002

CAPSULE, EXTENDED RELEASE;ORAL

MEPROSPAN

MEDPOINTE PHARM HLC

200MG

N011284 001

400MG

N011284 002

TABLET;ORAL

AMOSENE

FERNDAL LABS

400MG

A084030 001

BAMATE

ALRA

200MG

A080380 001

400MG

A080380 002

EQUANIL

WYETH AYERST

200MG

N010028 005

400MG

N010028 004

MEPRIAM

TEVA

400MG

N016069 001

MEPROBAMATE

ACELLA

400MG

A084153 001

BARR

600MG

A084230 001

CHARTWELL MOLECULAR

200MG

N014882 002

400MG

N014882 001

ELKINS SINN

200MG

N015426 002

400MG

N015426 001

HEATHER

400MG

N016928 003

600MG

A084329 001

IMPAX LABS

200MG

N014322 002

400MG

N014322 001

IVAX SUB TEVA PHARMS

200MG

N015438 001

400MG

N015438 002

600MG

A084181 001

LEDERLE

400MG

A086299 001

LEE KM

400MG

A089538 001

Nov 25, 1987

MALLARD

400MG

N015072 002

MK LABS

200MG

N014368 004

400MG

N014368 002

MYLAN

400MG

A083618 001

NEXGEN PHARMA INC

200MG

A084220 001

400MG

A084589 001

PARKE DAVIS

200MG

A084744 001

400MG

A084744 002

PERRIGO

200MG

A084546 001

400MG

A084547 001

PHARMAVITE

400MG

A084438 001

PUREPAC PHARM

200MG

A084804 001

400MG

A084804 002

PVT FORM

400MG

N014601 001

RISING

400MG

A080655 001

ROXANE

600MG

A084332 001

SANDOZ

200MG

N014547 002

400MG

N014547 001

SCHERER LABS

400MG

A083343 001

SOLVAY

200MG

A084435 001

STANLABS PHARM

200MG

N014474 002

400MG

N014474 004

SUN PHARM INDUSTRIES

200MG

A080699 001

400MG

A080699 002

TABLICAPS

400MG

A083494 001

TARO

200MG

A200998 001

May 23, 2011

400MG

A200998 002

May 23, 2011

USL PHARMA

200MG

A087825 001

Mar 18, 1982

400MG

A087826 001

Mar 18, 1982

VALEANT PHARM INTL

200MG

N015139 006

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

MEPROBAMATE

TABLET; ORAL

MEPROBAMATE

	400MG		N015139 005	
VANGARD	400MG		A088011 001	Jul 14, 1982
+ WATSON LABS	200MG		A083304 001	
	200MG		A085720 001	
+	400MG		A083308 001	
	400MG		A085721 001	
	600MG		A084274 001	
	600MG		A085719 001	
WEST WARD	200MG		N015417 003	
	400MG		N015417 002	
WHITEWORTH TOWN PLSN	200MG		A083830 001	
	400MG		A083442 001	
MILTOWN				
+ MEDPOINTE PHARM HLC	200MG **		N009698 004	
+	400MG **		N009698 002	
	600MG		A083919 001	
NEURAMATE				
HALSEY	200MG		N014359 002	
	400MG		N014359 001	
TRANMEP				
SOLVAY	400MG		A084369 001	
	400MG		N016249 001	

MEQUINOL; TRETINOIN

SOLUTION; TOPICAL

SOLAGE

ALMIRALL	2%;0.01%		N020922 001	Dec 10, 1999
----------	----------	--	-------------	--------------

MEROPENEM

INJECTABLE; INJECTION

MEROPENEM

DAEWOONG PHARM CO	500MG/VIAL		A204854 001	Dec 18, 2015
	1GM/VIAL		A204854 002	Dec 18, 2015
HOSPIRA INC	500MG/VIAL		A090940 001	Jun 22, 2010
	1GM/VIAL		A090940 002	Jun 22, 2010
SANDOZ	500MG/VIAL		A091201 001	Mar 29, 2011
	1GM/VIAL		A091201 002	Mar 29, 2011

MERSALYL SODIUM; THEOPHYLLINE

INJECTABLE; INJECTION

MERSALYL-THEOPHYLLINE

WATSON LABS	100MG/ML; 50MG/ML		A084875 001	
-------------	-------------------	--	-------------	--

MESALAMINE

CAPSULE, DELAYED RELEASE; ORAL

DELZICOL

+ ABBVIE	400MG		N204412 001	Feb 01, 2013
----------	-------	--	-------------	--------------

CAPSULE, EXTENDED RELEASE; ORAL

MESALAMINE

TEVA PHARMS USA	375MG		A209970 001	May 06, 2022
-----------------	-------	--	-------------	--------------

ENEMA; RECTAL

MESALAMINE

G AND W LABS INC	4GM/60ML		A076841 001	Sep 30, 2004
------------------	----------	--	-------------	--------------

SUPPOSITORY; RECTAL

CANASA

ABBVIE	500MG		N021252 001	Jan 05, 2001
--------	-------	--	-------------	--------------

MESALAMINE

AMNEAL	1GM		A210509 001	Jan 02, 2020
--------	-----	--	-------------	--------------

ROWASA

+ MEDA PHARMS	500MG **		N019919 001	Dec 18, 1990
---------------	----------	--	-------------	--------------

TABLET, DELAYED RELEASE; ORAL

ASACOL

APIL	400MG		N019651 001	Jan 31, 1992
------	-------	--	-------------	--------------

ASACOL HD

+ ABBVIE	800MG **		N021830 001	May 29, 2008
----------	----------	--	-------------	--------------

MESALAMINE

MYLAN	1.2GM		A203574 001	Nov 20, 2018
-------	-------	--	-------------	--------------

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

MESNA

INJECTABLE; INTRAVENOUS

MESNA

MYLAN INSTITUTIONAL	100MG/ML	A076488 001	Mar 08, 2012
RISING	100MG/ML	A203364 001	Jul 18, 2014
TEVA PHARMS USA	100MG/ML	A075764 001	Apr 27, 2001

MESORIDAZINE BESYLATE

CONCENTRATE; ORAL

SERENTIL

NOVARTIS	EQ 25MG BASE/ML	N016997 001	
----------	-----------------	-------------	--

INJECTABLE; INJECTION

SERENTIL

NOVARTIS	EQ 25MG BASE/ML	N016775 001	
----------	-----------------	-------------	--

TABLET; ORAL

SERENTIL

NOVARTIS	EQ 10MG BASE **	N016774 001	
	EQ 25MG BASE **	N016774 002	
	EQ 50MG BASE **	N016774 003	
	EQ 100MG BASE **	N016774 004	

MESTRANOL; NORETHINDRONE

TABLET; ORAL-20

NORINYL

ACTAVIS LABS UT INC	0.1MG; 2MG	N013625 004	
---------------------	------------	-------------	--

TABLET; ORAL-21

NORETHIN 1/50M-21

HERITAGE PHARMA	0.05MG; 1MG	A071539 001	Apr 12, 1988
-----------------	-------------	-------------	--------------

NORETHINDRONE AND MESTRANOL

WATSON LABS	0.05MG; 1MG	A070758 001	Jul 01, 1988
-------------	-------------	-------------	--------------

NORINYL 1+50 21-DAY

ACTAVIS LABS UT INC	0.05MG; 1MG	N013625 002	
---------------------	-------------	-------------	--

NORINYL 1+80 21-DAY

GD SEARLE LLC	0.08MG; 1MG	N016724 001	
---------------	-------------	-------------	--

ORTHO-NOVUM 1/50 21

ORTHO MCNEIL PHARM	0.05MG; 1MG	N012728 004	
--------------------	-------------	-------------	--

ORTHO-NOVUM 1/80 21

ORTHO MCNEIL PHARM	0.08MG; 1MG	N016715 001	
--------------------	-------------	-------------	--

ORTHO-NOVUM 10-21

ORTHO MCNEIL PHARM	0.06MG; 10MG	N012728 001	
--------------------	--------------	-------------	--

ORTHO-NOVUM 2-21

ORTHO MCNEIL PHARM	0.1MG; 2MG	N012728 005	
--------------------	------------	-------------	--

TABLET; ORAL-28

NORETHIN 1/50M-28

HERITAGE PHARMA	0.05MG; 1MG	A071540 001	Apr 12, 1988
-----------------	-------------	-------------	--------------

NORETHINDRONE AND MESTRANOL

WATSON LABS	0.05MG; 1MG	A070759 001	Jul 01, 1988
-------------	-------------	-------------	--------------

NORINYL 1+50 28-DAY

+ ACTAVIS LABS UT INC	0.05MG; 1MG	N016659 001	
-----------------------	-------------	-------------	--

NORINYL 1+80 28-DAY

GD SEARLE LLC	0.08MG; 1MG	N016725 001	
---------------	-------------	-------------	--

ORTHO-NOVUM 1/50 28

ORTHO MCNEIL JANSSEN	0.05MG; 1MG	N016709 001	
----------------------	-------------	-------------	--

ORTHO-NOVUM 1/80 28

ORTHO MCNEIL PHARM	0.08MG; 1MG	N016715 002	
--------------------	-------------	-------------	--

MESTRANOL; NORETHYNODREL

TABLET; ORAL

ENOVID

GD SEARLE LLC	0.075MG; 5MG	N010976 008	
	0.15MG; 9.85MG	N010976 005	

TABLET; ORAL-20

ENOVID

GD SEARLE LLC	0.075MG; 5MG	N010976 004	
---------------	--------------	-------------	--

ENOVID-E

GD SEARLE LLC	0.1MG; 2.5MG	N010976 006	
---------------	--------------	-------------	--

TABLET; ORAL-21

ENOVID-E 21

GD SEARLE LLC	0.1MG; 2.5MG	N010976 007	
---------------	--------------	-------------	--

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

METAPROTERENOL SULFATE

AEROSOL, METERED; INHALATION

ALUPENT

BOEHRINGER INGELHEIM 0.65MG/INH N016402 001

SOLUTION; INHALATION

ALUPENT

BOEHRINGER INGELHEIM 0.4% N018761 002 Oct 10, 1986

0.6% N018761 001 Jun 30, 1983

5% N017659 001

METAPROTERENOL SULFATE

APOTEX INC

0.4% A075402 001 Feb 28, 2001

0.6% A075403 001 Feb 28, 2001

ASTRAZENECA

0.4% A071275 001 Jul 27, 1988

0.6% A071018 001 Jul 27, 1988

DEY

0.33% A071806 001 Aug 05, 1988

0.5% A071805 001 Aug 05, 1988

5% A070805 001 Aug 17, 1987

MYLAN SPECIALITY LP

0.4% A071786 001 Aug 05, 1988

0.6% A070804 001 Aug 17, 1987

NEPHRON

0.4% A071855 001 Jul 14, 1988

0.6% A071726 001 Jul 14, 1988

WOCKHARDT

0.4% A075586 001 May 30, 2002

0.6% A075586 002 May 30, 2002

5% A072190 001 Jun 07, 1988

PROMETA

MURO

5% A073340 001 Mar 30, 1992

SYRUP; ORAL

ALUPENT

BOEHRINGER INGELHEIM 10MG/5ML N017571 001

METAPROTERENOL SULFATE

APOTEX INC

10MG/5ML A075235 001 Jan 27, 2000

COSETTE

10MG/5ML A072761 001 Feb 27, 1992

G AND W LABS INC

10MG/5ML A073034 001 Aug 30, 1991

MORTON GROVE

10MG/5ML A071656 001 Oct 13, 1987

WOCKHARDT

10MG/5ML A074702 001 Mar 24, 1997

PROMETA

MURO

10MG/5ML A072023 001 Sep 15, 1988

TABLET; ORAL

ALUPENT

BOEHRINGER INGELHEIM 10MG N015874 002

20MG N015874 001

METAPROTERENOL SULFATE

AM THERAP

10MG A072054 001 Jun 23, 1988

20MG A072055 001 Jun 23, 1988

HERITAGE PHARMA

10MG A072519 001 Mar 30, 1990

20MG A072520 001 Mar 30, 1990

STRIDES PHARMA

10MG A072024 001 Jun 28, 1988

20MG A072025 001 Jun 28, 1988

USL PHARMA

10MG A071013 001 Jan 25, 1988

20MG A071014 001 Jan 25, 1988

WATSON LABS

10MG A073013 001 Jan 31, 1991

20MG A072795 001 Jan 31, 1991

METARAMINOL BITARTRATE

INJECTABLE; INJECTION

ARAMINE

+ MERCK

EQ 10MG BASE/ML ** N009509 002 Dec 22, 1987

METARAMINOL BITARTRATE

ABRAXIS PHARM

EQ 10MG BASE/ML A080431 001

ELKINS SINN

EQ 10MG BASE/ML A083363 001

FRESENIUS KABI USA

EQ 10MG BASE/ML A080722 001

GD SEARLE LLC

EQ 10MG BASE/ML A086418 001

EQ 20MG BASE/ML A086418 002

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

METAXALONE

TABLET;ORAL

METAXALONE

INGENUS PHARMS LLC	800MG	A213836	001	Oct 21, 2020
+ PRIMUS PHARMS	640MG **	N022503	001	Jun 01, 2015
RISING	800MG	A208774	001	Sep 24, 2018
SKELAXIN				
+ KING PHARMS	400MG **	N013217	001	
+	800MG **	N013217	003	Aug 30, 2002

METFORMIN HYDROCHLORIDE

FOR SUSPENSION, EXTENDED RELEASE;ORAL

RIOMET ER

+ SUN PHARM	500MG/5ML	N212595	001	Aug 29, 2019
-------------	-----------	---------	-----	--------------

SOLUTION;ORAL

RIOMET

+ RANBAXY	500MG/5ML	N021591	001	Sep 11, 2003
-----------	-----------	---------	-----	--------------

TABLET;ORAL

GLUCOPHAGE

+ EMD SERONO INC	500MG **	N020357	001	Mar 03, 1995
+	625MG **	N020357	003	Nov 05, 1998
+	750MG **	N020357	004	Nov 05, 1998
+	850MG **	N020357	002	Mar 03, 1995
+	1GM **	N020357	005	Nov 05, 1998

METFORMIN HYDROCHLORIDE

AMNEAL PHARMS NY	500MG	A077880	001	Jun 05, 2006
	850MG	A077880	002	Jun 05, 2006
	1GM	A077880	003	Jun 05, 2006
BARR	500MG	A075971	001	Jan 25, 2002
	850MG	A075971	002	Jan 25, 2002
	1GM	A075971	003	Jan 25, 2002
HERITAGE PHARMA	500MG	A075978	001	Jan 25, 2002
	850MG	A075978	002	Jan 25, 2002
	1GM	A075978	003	Nov 05, 2002
INDICUS PHARMA	500MG	A079148	001	Nov 25, 2008
	850MG	A079148	002	Nov 25, 2008
	1GM	A079148	003	Nov 25, 2008
IPCA LABS LTD	500MG	A078422	001	Aug 06, 2007
	850MG	A078422	002	Aug 06, 2007
	1GM	A078422	003	Aug 06, 2007
IVAX SUB TEVA PHARMS	500MG	A075975	001	Jan 24, 2002
	625MG	A075975	004	Jan 24, 2002
	750MG	A075975	005	Jan 24, 2002
	850MG	A075975	002	Jan 24, 2002
	1GM	A075975	003	Jan 24, 2002
MACLEODS PHARMS LTD	500MG	A205330	001	Oct 31, 2017
	850MG	A205330	002	Oct 31, 2017
	1GM	A205330	003	Oct 31, 2017
NORVIUM BIOSCIENCE	500MG	A075969	001	Jan 29, 2002
	500MG	A075976	001	Jan 24, 2002
	850MG	A075969	002	Jan 29, 2002
	850MG	A075976	002	Jan 24, 2002
	1GM	A075969	003	Jan 29, 2002
	1GM	A075976	003	Jan 24, 2002
PROVIDENT PHARM	500MG	A077853	001	Jul 28, 2006
	850MG	A077853	002	Jul 28, 2006
	1GM	A077853	003	Jul 28, 2006
SANDOZ	500MG	A075985	001	Jan 25, 2002
	850MG	A075985	002	Jan 25, 2002
	1GM	A075985	003	Jan 25, 2002
SUN PHARM INDS INC	500MG	A075967	001	Jan 29, 2002
	850MG	A075967	002	Jan 29, 2002
	1GM	A075967	003	Jan 29, 2002
SUN PHARM INDUSTRIES	500MG	A076038	001	Feb 21, 2002
	850MG	A076038	002	Feb 21, 2002
	1GM	A076038	003	Feb 21, 2002
SUNSHINE	500MG	A208999	001	Oct 12, 2018
	850MG	A208999	002	Oct 12, 2018
	1GM	A208999	003	Oct 12, 2018
TEVA	500MG	A076328	001	Dec 16, 2002

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

METFORMIN HYDROCHLORIDE

TABLET;ORAL

METFORMIN HYDROCHLORIDE

	850MG		A076328	002	Dec 16, 2002
	1GM		A076328	003	Dec 16, 2002
TORRENT PHARMS	500MG		A077711	001	Jan 24, 2007
	850MG		A077711	002	Jan 24, 2007
	1GM		A077711	003	Jan 24, 2007
WATSON LABS	500MG		A075979	001	Jan 24, 2002
	850MG		A075979	002	Jan 24, 2002
	1GM		A075979	003	Jan 24, 2002
WATSON LABS FLORIDA	500MG		A075961	001	Jan 25, 2002
	850MG		A075961	002	Jan 25, 2002
	1GM		A075961	003	Jan 25, 2002

TABLET, EXTENDED RELEASE;ORAL

FORTAMET

+ ANDRX LABS LLC 500MG **

N021574 001 Apr 27, 2004

GLUCOPHAGE XR

+ EMD SERONO INC 500MG **

N021202 001 Oct 13, 2000

+ 750MG **

N021202 004 Apr 11, 2003

METFORMIN HYDROCHLORIDE

ACTAVIS ELIZABETH

500MG

A076450 001 Oct 01, 2004

750MG

A076878 001 Apr 13, 2005

ACTAVIS LABS FL INC

500MG

A076172 001 Jun 16, 2004

APOTEX

500MG

A076706 001 Dec 14, 2004

750MG

A076706 002 Dec 29, 2005

BARR

500MG

A076496 001 Nov 25, 2005

750MG

A076863 001 Oct 14, 2004

IMPAX LABS

500MG

A076249 001 Jul 30, 2004

750MG

A076985 001 Sep 13, 2005

IVAX SUB TEVA PHARMS

500MG

A076545 001 Dec 01, 2003

MACLEODS PHARMS LTD

1GM

A211163 001 Mar 13, 2024

NORVIUM BIOSCIENCE

500MG

A076650 001 Sep 13, 2005

750MG

A077113 001 Sep 08, 2005

RANBAXY LABS LTD

500MG

A076413 001 Jun 18, 2004

750MG

A077211 001 Jun 29, 2005

SANDOZ

500MG

A076223 001 Dec 14, 2004

SUN PHARM INDUSTRIES

500MG

A077124 001 Dec 21, 2005

TORRENT

500MG

A090014 001 Dec 30, 2009

TORRENT PHARMS LTD

750MG

A079226 001 Feb 18, 2010

UTOPIC PHARMS

500MG

A213394 001 Aug 03, 2021

1GM

A213394 002 Aug 03, 2021

WATSON LABS INC

500MG

A076818 001 Dec 14, 2004

METFORMIN HYDROCHLORIDE; PIOGLITAZONE HYDROCHLORIDE

TABLET;ORAL

ACTOPLUS MET

+ TAKEDA PHARMS USA 500MG;EQ 15MG BASE **

N021842 001 Aug 29, 2005

PIOGLITAZONE HYDROCHLORIDE AND METFORMIN HYDROCHLORIDE

CHARTWELL RX 500MG;EQ 15MG BASE

A091273 001 Apr 16, 2013

850MG;EQ 15MG BASE

A091273 002 Apr 16, 2013

NORVIUM BIOSCIENCE

500MG;EQ 15MG BASE

A090406 001 Feb 25, 2011

850MG;EQ 15MG BASE

A090406 002 Feb 25, 2011

TORRENT PHARMS LTD

500MG;EQ 15MG BASE

A202001 001 Feb 13, 2013

850MG;EQ 15MG BASE

A202001 002 Feb 13, 2013

TABLET, EXTENDED RELEASE;ORAL

ACTOPLUS MET XR

+ TAKEDA PHARMS USA 1GM;EQ 15MG BASE

N022024 001 May 12, 2009

+ 1GM;EQ 30MG BASE

N022024 002 May 12, 2009

METFORMIN HYDROCHLORIDE; REPAGLINIDE

TABLET;ORAL

PRANDIMET

+ NOVO NORDISK INC 500MG;1MG

N022386 001 Jun 23, 2008

+ 500MG;2MG

N022386 002 Jun 23, 2008

REPAGLINIDE AND METFORMIN HYDROCHLORIDE

LUPIN LTD 500MG;1MG

A200624 001 Jul 15, 2015

500MG;2MG

A200624 002 Jul 15, 2015

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

METFORMIN HYDROCHLORIDE; ROSIGLITAZONE MALEATE

TABLET; ORAL

AVANDAMET

+	SB PHARMCO	500MG;EQ 1MG BASE **	N021410 001	Oct 10, 2002
+		500MG;EQ 2MG BASE **	N021410 002	Oct 10, 2002
+		500MG;EQ 4MG BASE **	N021410 003	Oct 10, 2002
+		1GM;EQ 2MG BASE **	N021410 004	Aug 25, 2003
+		1GM;EQ 4MG BASE **	N021410 005	Aug 25, 2003

ROSIGLITAZONE MALEATE AND METFORMIN HYDROCHLORIDE

TEVA

		500MG;EQ 2MG BASE	A077337 001	May 07, 2014
		500MG;EQ 1MG BASE	A077337 005	May 19, 2017
		500MG;EQ 4MG BASE	A077337 002	May 07, 2014
		1GM;EQ 4MG BASE	A077337 004	May 07, 2014
		1GM;EQ 2MG BASE	A077337 003	May 07, 2014

METFORMIN HYDROCHLORIDE; SAXAGLIPTIN HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL

KOMBIGLYZE XR

+	ASTRAZENECA AB	500MG;EQ 5MG BASE **	N200678 001	Nov 05, 2010
+		1GM;EQ 2.5MG BASE **	N200678 003	Nov 05, 2010
+		1GM;EQ 5MG BASE **	N200678 002	Nov 05, 2010

METHACHOLINE CHLORIDE

FOR SOLUTION; INHALATION

PROVOCHOLINE

+	METHAPHARM	1600MG/VIAL	N019193 002	Aug 29, 2016
---	------------	-------------	-------------	--------------

METHACYCLINE HYDROCHLORIDE

CAPSULE; ORAL

RONDONMYCIN

	MEDPOINTE PHARM HLC	EQ 140MG BASE	A060641 001	
		EQ 280MG BASE	A060641 002	

SYRUP; ORAL

RONDONMYCIN

	MEDPOINTE PHARM HLC	EQ 70MG BASE/5ML	A060641 003	
--	---------------------	------------------	-------------	--

METHADONE HYDROCHLORIDE

CONCENTRATE; ORAL

METHADONE HYDROCHLORIDE

	LANNETT CO INC	10MG/ML	A212094 001	Mar 03, 2021
--	----------------	---------	-------------	--------------

POWDER; FOR RX COMPOUNDING

METHADONE HYDROCHLORIDE

	MALLINCKRODT INC	50GM/BOT	N006383 002	
		100GM/BOT	N006383 003	
		500GM/BOT	N006383 004	

SOLUTION; ORAL

METHADONE HYDROCHLORIDE

	VISTAPHARM LLC	5MG/5ML	A090707 001	Jun 30, 2010
		10MG/5ML	A090707 002	Jun 30, 2010

SYRUP; ORAL

DOLOPHINE HYDROCHLORIDE

	HIKMA	10MG/30ML	N006134 004	
--	-------	-----------	-------------	--

TABLET; ORAL

DOLOPHINE HYDROCHLORIDE

+	HIKMA	5MG **	N006134 002	
+		10MG **	N006134 010	

METHADONE HYDROCHLORIDE

	ROXANE	40MG	A074081 001	Apr 28, 1995
	SUN PHARM INDUSTRIES	5MG	A208305 001	Mar 30, 2018
		10MG	A208305 002	Mar 30, 2018
	VISTAPHARM LLC	5MG	A040241 001	May 29, 1998

METHADOSE

	SPECGX LLC	5MG	A040050 001	Apr 15, 1993
		10MG	A040050 002	Apr 15, 1993

TABLET, DISPERSIBLE; ORAL

WESTADONE

	SANDOZ	2.5MG	N017108 001	
--	--------	-------	-------------	--

TABLET, EFFERVESCENT; ORAL

WESTADONE

	SANDOZ	5MG	N017108 002	
		10MG	N017108 003	

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

METHADONE HYDROCHLORIDETABLET, EFFERVESCENT;ORAL
WESTADONE

40MG

N017108 004

METHAMPHETAMINE HYDROCHLORIDE

TABLET;ORAL

METHAMPEX

TEVA 10MG

A083889 001

METHAMPHETAMINE HYDROCHLORIDE

ABLE 5MG

A040529 001 Feb 25, 2004

REXAR 5MG

A084931 001

10MG

A084931 002

TEVA 5MG

A086359 001

TABLET, EXTENDED RELEASE;ORAL

DESOXYN

AJENAT PHARMS 5MG

N005378 004

10MG

N005378 003

15MG

N005378 005

METHANTHELINE BROMIDE

TABLET;ORAL

BANTHINE

SHIRE 50MG

N007390 001

METHARBITAL

TABLET;ORAL

GEMONIL

ABBVIE 100MG

N008322 001

METHAZOLAMIDE

TABLET;ORAL

METHAZOLAMIDE

APPLIED ANAL 25MG

A040011 001 Jul 17, 1997

50MG

A040011 002 Jul 17, 1997

ATHEM 25MG

A040102 001 Aug 28, 1996

50MG

A040102 002 Aug 28, 1996

NEPTAZANE

+ LEDERLE 25MG **

N011721 002 Nov 25, 1991

+ 50MG **

N011721 001

METHDILAZINE

TABLET, CHEWABLE;ORAL

TACARYL

WESTWOOD SQUIBB 3.6MG

N011950 009

METHDILAZINE HYDROCHLORIDE

SYRUP;ORAL

METHDILAZINE HYDROCHLORIDE

ALPHARMA US PHARMS 4MG/5ML

A087122 001

TACARYL

WESTWOOD SQUIBB 4MG/5ML

N011950 007

TABLET;ORAL

TACARYL

WESTWOOD SQUIBB 8MG

N011950 006

METHENAMINE HIPPURATE

TABLET;ORAL

METHENAMINE HIPPURATE

IMPAX LABS INC 1GM

A076411 001 Jun 20, 2003

METHICILLIN SODIUM

INJECTABLE; INJECTION

STAPHCILLIN

APOTHECON EQ 900MG BASE/VIAL

A061449 001

EQ 900MG BASE/VIAL

N050117 001

EQ 3.6GM BASE/VIAL

A061449 002

EQ 3.6GM BASE/VIAL

N050117 002

EQ 5.4GM BASE/VIAL

A061449 003

EQ 5.4GM BASE/VIAL

N050117 003

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

METHIMAZOLE

TABLET; ORAL

METHIMAZOLE

DISCOVERY THERAP	15MG	A040619	003	Jul 12, 2005
MYLAN	20MG	A040350	003	Jun 07, 2001
QINGDAO BAHEAL PHARM	20MG	A040547	004	Feb 18, 2005
SUN PHARM INDS INC	5MG	A040870	001	Sep 25, 2007
	10MG	A040870	002	Sep 25, 2007

TAPAZOLE

+ KING PHARMS	5MG **	N007517	002	
+ KING PHARMS LLC	10MG **	N007517	004	
	5MG	A040320	001	Mar 31, 2000
	10MG	A040320	002	Mar 31, 2000

METHIXENE HYDROCHLORIDE

TABLET; ORAL

TREST

NOVARTIS	1MG	N013420	001	
----------	-----	---------	-----	--

METHOCARBAMOL

INJECTABLE; INJECTION

METHOCARBAMOL

DR REDDYS	100MG/ML	A086459	001	
MARSAM PHARMS LLC	100MG/ML	A089849	001	Dec 27, 1991

SOLUTION; IM-IV

METHOCARBAMOL

BAXTER HLTHCARE CORP	1GM/10ML (100MG/ML)	A215065	001	Jul 14, 2022
NAVINTA LLC	1GM/10ML (100MG/ML)	A206071	001	Nov 24, 2017
NORVIUM BIOSCIENCE	1GM/10ML (100MG/ML)	A204404	001	Dec 05, 2014

TABLET; ORAL

DELAXIN

FERNDALE LABS	500MG	A085454	001	
---------------	-------	---------	-----	--

FORBAXIN

FOREST LABS	750MG	A085136	001	
-------------	-------	---------	-----	--

METHOCARBAMOL

ABLE	500MG	A040413	001	Mar 17, 2003
	750MG	A040413	002	Mar 17, 2003
AIPING PHARM INC	500MG	A084616	001	
	750MG	A084615	001	
AM THERAP	500MG	A089417	001	Feb 11, 1987
	750MG	A089418	001	Feb 11, 1987
ANI PHARMS	500MG	A084277	001	
	750MG	A084276	002	
ASCOT	500MG	A087660	001	Oct 27, 1982
	750MG	A087661	001	Oct 27, 1982
CHARTWELL MOLECULAR	500MG	A084756	002	Mar 31, 2003
	750MG	A084756	001	
CLONMEL HLTHCARE	500MG	A085961	001	
	750MG	A085963	001	
HEATHER	500MG	A084675	001	
	750MG	A084924	001	
HIKMA INTL PHARMS	500MG	A085159	001	
	750MG	A085123	001	
IMPAX LABS	500MG	A084927	001	
	750MG	A084928	001	
INWOOD LABS	500MG	A085137	001	
IVAX SUB TEVA PHARMS	500MG	A084648	001	
	750MG	A084649	001	
KV PHARM	500MG	A085660	001	
	750MG	A085658	001	
MYLAN	500MG	A084259	001	
	750MG	A084323	001	
NYLOS	750MG	A085033	001	
PIONEER PHARMS	500MG	A088731	001	Dec 13, 1985
	750MG	A089082	001	Dec 13, 1985
PURACAP PHARM	500MG	A084231	002	
	750MG	A084471	001	
PUREPAC PHARM	500MG	A085718	001	
	750MG	A085718	002	
ROXANE	500MG	A088646	001	Feb 29, 1984

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

METHOCARBAMOL

TABLET; ORAL

METHOCARBAMOL

	750MG	A088647 001	Feb 29, 1984
SANDOZ	500MG	A087283 001	
	750MG	A087282 001	
SOLVAY	500MG	A084448 001	
	750MG	A084449 001	
SUN PHARM INDUSTRIES	500MG	A084488 001	
	750MG	A084486 001	
SUPERPHARM	500MG	A087589 001	Jan 22, 1982
	750MG	A087590 001	Jan 22, 1982
TABLICAPS	500MG	A084846 001	
UPSHER SMITH	500MG	A087453 001	
	750MG	A087454 001	
WATSON LABS	500MG	A083605 001	
	500MG	A085180 001	
	750MG	A083605 002	
	750MG	A085192 001	
ROBAXIN			
+ ENDO OPERATIONS	500MG **	N011011 004	
ROBAXIN-750			
+ ENDO OPERATIONS	750MG **	N011011 006	

METHOHEXITAL SODIUM

INJECTABLE; INJECTION

BREVITAL SODIUM

ENDO OPERATIONS	200MG/VIAL	N011559 004	Dec 21, 2012
+	2.5GM/VIAL	N011559 002	
	5GM/VIAL	N011559 003	

METHOTREXATE

SOLUTION; INTRAVENOUS

METHOTREXATE

+ ACCORD HLTHCARE	5GM/50ML (100MG/ML)	N214121 001	Aug 24, 2020
-------------------	---------------------	-------------	--------------

SOLUTION; SUBCUTANEOUS

OTREXUP

+ OTTER PHARMS	7.5MG/0.4ML (7.5MG/0.4ML)	N204824 005	Nov 07, 2014
----------------	---------------------------	-------------	--------------

OTREXUP PFS

+ OTTER PHARMS	10MG/0.4ML (10MG/0.4ML)	N204824 009	May 31, 2017
+	15MG/0.6ML (15MG/0.6ML)	N204824 010	May 31, 2017
+	17.5MG/0.7ML (17.5MG/0.7ML)	N204824 011	May 31, 2017
+	20MG/0.8ML (20MG/0.8ML)	N204824 012	May 31, 2017
+	22.5MG/0.9ML (22.5MG/0.9ML)	N204824 013	May 31, 2017
+	25MG/ML (25MG/ML)	N204824 014	May 31, 2017

RASUVO

+ MEDEXUS	27.5MG/0.55ML (27.5MG/0.55ML)	N205776 009	Jul 10, 2014
-----------	-------------------------------	-------------	--------------

REDITREX

+ NORDIC GRP	7.5MG/0.3ML (7.5MG/0.3ML)	N210737 001	Nov 27, 2019
+	10MG/0.4ML (10MG/0.4ML)	N210737 002	Nov 27, 2019
+	12.5MG/0.5ML (12.5MG/0.5ML)	N210737 003	Nov 27, 2019
+	15MG/0.6ML (15MG/0.6ML)	N210737 004	Nov 27, 2019
+	17.5MG/0.7ML (17.5MG/0.7ML)	N210737 005	Nov 27, 2019
+	20MG/0.8ML (20MG/0.8ML)	N210737 006	Nov 27, 2019
+	22.5MG/ML (22.5MG/ML)	N210737 007	Nov 27, 2019
+	25MG/1ML (25MG/1ML)	N210737 008	Nov 27, 2019

METHOTREXATE SODIUM

INJECTABLE; INJECTION

ABITREXATE

ABIC	EQ 25MG BASE/ML	A089161 001	Mar 10, 1987
	EQ 50MG BASE/VIAL	A089354 001	Jul 17, 1987
	EQ 100MG BASE/VIAL	A089355 001	Jul 17, 1987
	EQ 250MG BASE/VIAL	A089356 001	Jul 17, 1987

FOLEX

+ PHARMACIA AND UPJOHN	EQ 25MG BASE/VIAL	A087695 001	Apr 08, 1983
	EQ 50MG BASE/VIAL	A087695 002	Apr 08, 1983
+	EQ 100MG BASE/VIAL	A087695 003	Apr 08, 1983
	EQ 250MG BASE/VIAL	A088954 001	Oct 24, 1985

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

METHOTREXATE SODIUM

INJECTABLE; INJECTION

FOLEX PFS

PHARMACIA AND UPJOHN	EQ 25MG BASE/ML	A081242	001	Aug 23, 1991
	EQ 25MG BASE/ML	A089180	001	Jan 03, 1986

METHOTREXATE LPF

+ HOSPIRA	EQ 25MG BASE/ML	N011719	007	Mar 31, 1982
-----------	-----------------	---------	-----	--------------

METHOTREXATE PRESERVATIVE FREE

FRESENIUS KABI USA	EQ 25MG BASE/ML	A040265	001	Feb 26, 1999
+ HOSPIRA	EQ 20MG BASE/2ML (EQ 10MG BASE/ML)	N011719	014	Apr 13, 2005
+ HOSPIRA	EQ 500MG BASE/20ML (EQ 25MG BASE/ML) **	N011719	013	Apr 13, 2005
+ HOSPIRA	EQ 2.5GM BASE/100ML (EQ 25MG BASE/ML)	N011719	011	Apr 13, 2005

METHOTREXATE SODIUM

ABRAXIS PHARM

	EQ 2.5MG BASE/ML	A089323	001	Jun 13, 1986
	EQ 20MG BASE/VIAL	A088935	001	Oct 11, 1985
	EQ 25MG BASE/ML	A089263	001	Jun 13, 1986
	EQ 25MG BASE/ML	A089322	001	Jun 13, 1986
	EQ 50MG BASE/VIAL	A088936	001	Oct 11, 1985
	EQ 100MG BASE/VIAL	A088937	001	Oct 11, 1985

FRESENIUS KABI USA	EQ 250MG BASE/10ML (EQ 25MG BASE/ML)	A040263	002	Feb 26, 1999
--------------------	--------------------------------------	---------	-----	--------------

+ HOSPIRA	EQ 2.5MG BASE/ML	N011719	004	
+ HOSPIRA	EQ 20MG BASE/VIAL	N011719	001	
+ HOSPIRA	EQ 25MG BASE/ML	N011719	005	
+ HOSPIRA	EQ 50MG BASE/VIAL	N011719	003	
+ HOSPIRA	EQ 100MG BASE/VIAL	N011719	006	

NORBROOK	EQ 25MG BASE/ML	A088648	001	May 09, 1986
----------	-----------------	---------	-----	--------------

PHARMACHEMIE USA	EQ 25MG BASE/ML	A089158	001	Jul 08, 1988
------------------	-----------------	---------	-----	--------------

METHOTREXATE SODIUM PRESERVATIVE FREE

EUGIA PHARMA SPECLTS	EQ 50MG BASE/2ML (EQ 25MG BASE/ML)	A201529	001	Mar 29, 2012
	EQ 100MG BASE/4ML (EQ 25MG BASE/ML)	A201529	002	Mar 29, 2012
	EQ 200MG BASE/8ML (EQ 25MG BASE/ML)	A201529	003	Mar 29, 2012
	EQ 250MG BASE/10ML (EQ 25MG BASE/ML)	A201529	004	Mar 29, 2012
EXTROVIS	EQ 1GM BASE/40ML (EQ 25MG BASE/ML)	A201530	001	Mar 29, 2012

+ HOSPIRA	EQ 1GM BASE/VIAL	N011719	009	Apr 07, 1988
-----------	------------------	---------	-----	--------------

MEXATE

+ BRISTOL	EQ 20MG BASE/VIAL	A086358	001	
+ BRISTOL	EQ 50MG BASE/VIAL	A086358	002	
+ BRISTOL	EQ 100MG BASE/VIAL	A086358	003	
+ BRISTOL	EQ 250MG BASE/VIAL	A086358	004	

MEXATE-AQ

+ BRISTOL MYERS	EQ 25MG BASE/ML	A088760	001	Feb 14, 1985
-----------------	-----------------	---------	-----	--------------

MEXATE-AQ PRESERVED

BRISTOL MYERS SQUIBB	EQ 25MG BASE/ML	A089887	001	Apr 14, 1989
----------------------	-----------------	---------	-----	--------------

TABLET; ORAL

METHOTREXATE SODIUM

AMNEAL PHARMS	EQ 2.5MG BASE	A210040	001	Dec 22, 2017
DURAMED PHARMS BARR	EQ 2.5MG BASE	A040233	001	Jun 17, 1999
LOTUS PHARM CO LTD	EQ 2.5MG BASE	A209787	001	Apr 23, 2021
+ STRIDES PHARMA	EQ 2.5MG BASE **	N008085	002	

METHOXAMINE HYDROCHLORIDE

INJECTABLE; INJECTION

VASOXYL

GLAXOSMITHKLINE	10MG/ML	N006772	002	
	20MG/ML	N006772	001	

METHOXSALEN

CAPSULE; ORAL

8-MOP

+ VALEANT PHARM INTL	10MG	N009048	001	
----------------------	------	---------	-----	--

METHOXSALEN

ACTAVIS INC	10MG	A202603	001	Jun 09, 2015
ANI PHARMS	10MG	A087781	001	Jun 08, 1982

LOTION; TOPICAL

OXSORALEN

+ VALEANT PHARM INTL	1%	N009048	002	
----------------------	----	---------	-----	--

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

METHSCOPOLAMINE BROMIDE

TABLET; ORAL

METHSCOPOLAMINE BROMIDE

CHARTWELL RX	2.5MG	A040624 001	Dec 28, 2006
	5MG	A040624 002	Dec 28, 2006
PVT FORM	2.5MG	A080970 001	

PAMINE

FOUGERA PHARMS	2.5MG **	N008848 001	
----------------	----------	-------------	--

PAMINE FORTE

FOUGERA PHARMS	5MG **	N008848 002	Mar 25, 2003
----------------	--------	-------------	--------------

METHSUXIMIDE

CAPSULE; ORAL

CELONTIN

+ PARKE DAVIS	150MG	N010596 007	
---------------	-------	-------------	--

METHYCLOTHIAZIDE

TABLET; ORAL

AQUATENSEN

MEDPOINTE PHARM HLC	5MG	N017364 001	
---------------------	-----	-------------	--

ENDURON

+ ABBVIE	2.5MG **	N012524 001	
----------	----------	-------------	--

+	5MG **	N012524 004	
---	--------	-------------	--

METHYCLOTHIAZIDE

CHARTWELL RX	2.5MG	A089835 001	Aug 18, 1988
	5MG	A089837 001	Aug 18, 1988
IVAX PHARMS	2.5MG	A087913 001	Jun 03, 1982
	5MG	A087786 001	May 18, 1982
MYLAN	2.5MG	A087671 001	Aug 17, 1982
NORVIUM BIOSCIENCE	5MG	A087672 001	Aug 17, 1982
PAR PHARM	2.5MG	A089135 001	Feb 12, 1986
	5MG	A089136 001	Feb 12, 1986
USL PHARMA	5MG	A088745 001	Mar 21, 1985
WATSON LABS	2.5MG	A085487 001	Mar 11, 1982
	2.5MG	A088750 001	Sep 06, 1984
	5MG	A085476 001	Mar 11, 1982
	5MG	A088724 001	Sep 06, 1984

METHYCLOTHIAZIDE; PARGYLINE HYDROCHLORIDE

TABLET; ORAL

EUTRON

ABBOTT	5MG;25MG	N016047 001	
--------	----------	-------------	--

METHYCLOTHIAZIDE; RESERPINE

TABLET; ORAL

DIUTENSEN-R

MEDPOINTE PHARM HLC	2.5MG;0.1MG	N012708 005	
---------------------	-------------	-------------	--

METHYL AMINOLEVULINATE HYDROCHLORIDE

CREAM; TOPICAL

METVIXIA

GALDERMA LABS LP	EQ 16.8% BASE	N021415 001	Jul 27, 2004
------------------	---------------	-------------	--------------

METHYLDOPA

SUSPENSION; ORAL

ALDOMET

MERCK	250MG/5ML	N018389 001	
-------	-----------	-------------	--

TABLET; ORAL

ALDOMET

+ MERCK	125MG **	N013400 003	
---------	----------	-------------	--

+	250MG **	N013400 001	
---	----------	-------------	--

+	500MG **	N013400 002	
---	----------	-------------	--

METHYLDOPA

ACCORD HLTHCARE	125MG	A070070 003	Oct 15, 1985
	250MG	A070084 001	Oct 15, 1985
	500MG	A070085 001	Oct 15, 1985
CHARTWELL RX	125MG	A071700 001	Mar 02, 1988
	250MG	N018934 001	Jun 29, 1984
	500MG	N018934 002	Jun 29, 1984
DURAMED PHARMS BARR	250MG	A071006 001	Dec 16, 1986
	500MG	A071009 001	Dec 16, 1986
HALSEY	125MG	A071751 001	Mar 28, 1988

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

METHYLDOPATABLET; ORAL
METHYLDOPA

	250MG	A071752 001	Mar 28, 1988
	500MG	A071753 001	Mar 28, 1988
HERITAGE PHARMA	250MG	A070098 001	Feb 20, 1986
	500MG	A070343 001	Feb 20, 1986
PARKE DAVIS	125MG	A070331 001	Apr 15, 1986
	250MG	A070332 001	Apr 15, 1986
	500MG	A070333 001	Apr 15, 1986
PLIVA	125MG	A072126 001	Jul 07, 1988
	250MG	A072127 001	Jul 07, 1988
	500MG	A072128 001	Jul 07, 1988
PUREPAC PHARM	125MG	A070749 001	Feb 07, 1986
	250MG	A070750 001	Feb 07, 1986
	500MG	A070452 001	Feb 07, 1986
ROXANE	125MG	A070192 001	Apr 25, 1986
	250MG	A070193 001	Apr 25, 1986
	500MG	A070194 001	Apr 25, 1986
STRIDES PHARMA	125MG	A070535 001	Jan 02, 1987
	250MG	A070536 001	Jan 02, 1987
	500MG	A070537 001	Jan 02, 1987
SUN PHARM INDUSTRIES	125MG	A070073 001	Oct 09, 1986
	250MG	A070060 001	Oct 09, 1986
	500MG	A070074 001	Oct 09, 1986
SUPERPHARM	250MG	A070669 001	Jun 23, 1989
	500MG	A070670 001	Jun 23, 1989
TEVA	125MG	A071105 001	Dec 05, 1986
	250MG	A071106 001	Dec 05, 1986
	500MG	A071067 001	Dec 05, 1986
WATSON LABS	125MG	A070245 001	Feb 25, 1986
	125MG	A070260 001	Jun 24, 1985
	250MG	A070246 001	Feb 25, 1986
	250MG	A070261 001	Jun 24, 1985
	250MG	A070703 001	Jun 06, 1986
	500MG	A070247 001	Feb 25, 1986
	500MG	A070262 001	Jun 24, 1985
	500MG	A070625 001	Jun 06, 1986

METHYLDOPATE HYDROCHLORIDE

INJECTABLE; INJECTION

ALDOMET

+ MERCK 50MG/ML **

N013401 001

METHYLDOPATE HYDROCHLORIDE

ABRAXIS PHARM	50MG/ML	A070652 001	Jun 03, 1986
AM REGENT	50MG/ML	A071279 001	Oct 02, 1987
BAXTER HLTHCARE	50MG/ML	A070291 001	Jul 01, 1986
HOSPIRA	50MG/ML	A070691 001	Jun 19, 1987
	50MG/ML	A070698 001	Jun 15, 1987
	50MG/ML	A070699 001	Jun 15, 1987
	50MG/ML	A070849 001	Jun 19, 1987
MARSAM PHARMS LLC	50MG/ML	A071812 001	Dec 22, 1987
SMITH AND NEPHEW	50MG/ML	A070841 001	Jan 02, 1987
TEVA PARENTERAL	50MG/ML	A072974 001	Nov 22, 1991

METHYLERGONOVINE MALEATE

INJECTABLE; INJECTION

METHERGINE

+ EDISON THERAPS LLC 0.2MG/ML

N006035 004

TABLET; ORAL

METHERGINE

+ EDISON THERAPS LLC 0.2MG **

N006035 003

METHYLPHENIDATE

TABLET, ORALLY DISINTEGRATING, EXTENDED RELEASE; ORAL

COTEMPLA XR-ODT

+ NEOS THERAPS INC 34.6MG

N205489 004 Aug 19, 2024

METHYLPHENIDATE

ACTAVIS ELIZABETH	8.6MG	A210924 001	Jun 19, 2020
	17.3MG	A210924 002	Jun 19, 2020
	25.9MG	A210924 003	Jun 19, 2020

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

METHYLPHENIDATE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE;ORAL

ADHANSIA XR

+	PURDUE PHARMA LP	25MG	N212038	001	Feb 27, 2019
+		35MG	N212038	002	Feb 27, 2019
+		45MG	N212038	003	Feb 27, 2019
+		55MG	N212038	004	Feb 27, 2019
+		70MG	N212038	005	Feb 27, 2019
+		85MG	N212038	006	Feb 27, 2019

METHYLPHENIDATE HYDROCHLORIDE

BARR LABS INC

		10MG	A079031	004	Oct 15, 2014
		20MG	A079031	001	Jul 13, 2012
		30MG	A079031	002	Jul 13, 2012
		40MG	A079031	003	Jul 13, 2012
	TEVA PHARMS	40MG	A078873	001	Jul 19, 2012
		50MG	A078873	002	Jul 19, 2012
		60MG	A078873	003	Jul 19, 2012

RITALIN LA

+	SANDOZ	60MG **	N021284	005	Oct 27, 2014
---	--------	---------	---------	-----	--------------

SOLUTION;ORAL

METHYLPHENIDATE HYDROCHLORIDE

ALLIED

		5MG/5ML	A201466	001	Nov 12, 2013
		10MG/5ML	A201466	002	Nov 12, 2013

CHARTWELL MOLECULAR

		5MG/5ML	A207414	001	Dec 16, 2020
		10MG/5ML	A207414	002	Dec 16, 2020

TABLET;ORAL

METHYLPHENIDATE HYDROCHLORIDE

ABLE

		5MG	A040404	001	Mar 29, 2001
		10MG	A040404	002	Mar 29, 2001
		20MG	A040404	003	Mar 29, 2001

ACTAVIS ELIZABETH

		5MG	A040321	001	Feb 05, 2002
		10MG	A040321	002	Feb 05, 2002
		20MG	A040321	003	Feb 05, 2002

ALVOGEN

		5MG	A206840	001	Sep 15, 2016
		10MG	A206840	002	Sep 15, 2016
		20MG	A206840	003	Sep 15, 2016

AUROLIFE PHARMA LLC

		5MG	A209276	001	Oct 25, 2018
		10MG	A209276	002	Oct 25, 2018
		20MG	A209276	003	Oct 25, 2018

LANNETT CO INC

		5MG	A086429	001	
		10MG	A085799	001	
		20MG	A086428	001	

NOSTRUM LABS INC

		5MG	A207587	001	Mar 03, 2017
		10MG	A207587	002	Mar 03, 2017
		20MG	A207587	003	Mar 03, 2017

WATSON LABS

		5MG	A040220	001	Aug 29, 1997
		10MG	A040220	002	Aug 29, 1997
		20MG	A040220	003	Aug 29, 1997

TABLET, CHEWABLE;ORAL

METHYLIN

+	SPECGX LLC	2.5MG **	N021475	001	Apr 15, 2003
+		5MG **	N021475	002	Apr 15, 2003
+		10MG **	N021475	003	Apr 15, 2003

METHYLPHENIDATE HYDROCHLORIDE

NOSTRUM LABS INC

		2.5MG	A204954	001	Jan 26, 2017
		5MG	A204954	002	Jan 26, 2017
		10MG	A204954	003	Jan 26, 2017

NOVEL LABS INC

		2.5MG	A204115	001	Feb 25, 2015
		5MG	A204115	002	Feb 25, 2015
		10MG	A204115	003	Feb 25, 2015

TABLET, EXTENDED RELEASE;ORAL

METADATE ER

LANNETT CO INC

		10MG	A040306	001	Oct 20, 1999
		20MG	A089601	001	Jun 01, 1988

METHYLPHENIDATE HYDROCHLORIDE

ABLE

		20MG	A076032	001	May 09, 2001
--	--	------	---------	-----	--------------

ALVOGEN

		10MG	A204772	001	Feb 29, 2016
		18MG	A210818	001	Nov 30, 2018
		20MG	A204772	002	Feb 29, 2016

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

METHYLPHENIDATE HYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL

METHYLPHENIDATE HYDROCHLORIDE

	27MG	A210818 002	Nov 30, 2018
	36MG	A210818 003	Nov 30, 2018
	54MG	A210818 004	Nov 30, 2018
AMNEAL PHARMS	18MG	A207515 001	Feb 01, 2018
	27MG	A207515 002	Feb 01, 2018
	36MG	A207515 003	Feb 01, 2018
	54MG	A207515 004	Feb 01, 2018
ANI PHARMS	18MG	A208607 001	Jul 14, 2017
	27MG	A208607 002	Jul 14, 2017
	36MG	A208607 003	Jul 14, 2017
	54MG	A208607 004	Jul 14, 2017
HERITAGE PHARMA	20MG	A075450 001	Dec 21, 2001
RHODES PHARMS	18MG	A214111 001	May 31, 2022
	27MG	A214111 002	May 31, 2022
	36MG	A214111 003	May 31, 2022
	54MG	A214111 004	May 31, 2022
STRIDES PHARMA	36MG	A204659 001	Jul 15, 2019
	54MG	A204659 002	Jul 15, 2019
WATSON LABS	20MG	A040410 001	Feb 09, 2001
RITALIN-SR			
+ NOVARTIS	20MG **	N018029 001	Mar 30, 1982

METHYLPREDNISOLONE

TABLET;ORAL

MEDROL

PFIZER	24MG	N011153 005	
METHYLPREDNISOLONE			
AMNEAL	4MG	A207481 001	Sep 21, 2021
	8MG	A207481 002	Sep 21, 2021
	16MG	A207481 003	Sep 21, 2021
	32MG	A207481 004	Sep 21, 2021
CHARTWELL RX	2MG	A209097 001	Feb 22, 2019
	4MG	A209097 002	Feb 22, 2019
	8MG	A209097 003	Feb 22, 2019
	16MG	A209097 004	Feb 22, 2019
	32MG	A209097 005	Feb 22, 2019
DURAMED PHARMS BARR	4MG	A088497 001	Feb 21, 1984
HEATHER	4MG	A085650 001	
INVATECH	4MG	A087341 001	
NOVAST LABS	4MG	A210985 001	Jan 09, 2019
PAR PHARM	16MG	A089207 001	Apr 25, 1988
	24MG	A089208 001	Apr 25, 1988
	32MG	A089209 001	Apr 25, 1988
PRAXGEN	4MG	A212262 001	Jun 27, 2019
WATSON LABS	4MG	A086161 001	Feb 09, 1982
	16MG	A086159 001	Feb 09, 1982

METHYLPREDNISOLONE ACETATE

ENEMA;RECTAL

MEDROL

PHARMACIA AND UPJOHN	40MG/BOT	N018102 001	
INJECTABLE; INJECTION			
M-PREDROL			
BEL MAR	40MG/ML	A086666 001	
	80MG/ML	A087135 001	
METHYLPREDNISOLONE ACETATE			
EPIC PHARMA LLC	40MG/ML	A086903 001	Oct 20, 1982
	80MG/ML	A086903 002	Oct 20, 1982
SAGENT PHARMS INC	20MG/ML	A201835 001	Jun 27, 2018
TEVA PHARMS USA	40MG/ML	A040620 001	Oct 27, 2006
	80MG/ML	A040620 002	Oct 27, 2006
WATSON LABS	20MG/ML	A085597 001	
	20MG/ML	A087248 001	
	40MG/ML	A085374 001	
	40MG/ML	A085600 001	
	80MG/ML	A085595 001	
	80MG/ML	A086507 001	

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

METHYLPREDNISOLONE ACETATE

OINTMENT; TOPICAL

MEDROL ACETATE

PHARMACIA AND UPJOHN	0.25%	N012421	001
	1%	N012421	002

METHYLPREDNISOLONE ACETATE; NEOMYCIN SULFATE

CREAM; TOPICAL

NEO-MEDROL ACETATE

PHARMACIA AND UPJOHN	0.25%;EQ 3.5MG BASE/GM	A060611	002
	1%;EQ 3.5MG BASE/GM	A060611	001

METHYLPREDNISOLONE SODIUM SUCCINATE

INJECTABLE; INJECTION

A-METHAPRED

ABBOTT	EQ 40MG BASE/VIAL	A089573	001	Feb 22, 1991
	EQ 125MG BASE/VIAL	A089574	001	Feb 22, 1991
	EQ 500MG BASE/VIAL	A089575	001	Feb 22, 1991
HOSPIRA	EQ 1GM BASE/VIAL	A089576	001	Feb 22, 1991
	EQ 40MG BASE/VIAL	A040664	001	Dec 20, 2005
	EQ 40MG BASE/VIAL	A085853	001	
HOSPIRA	EQ 125MG BASE/VIAL	A040665	001	Dec 20, 2005
	EQ 125MG BASE/VIAL	A085855	001	
	EQ 500MG BASE/VIAL	A085854	001	
	EQ 500MG BASE/VIAL	A089173	001	Aug 18, 1987
	EQ 1GM BASE/VIAL	A085852	001	
	EQ 1GM BASE/VIAL	A089174	001	Aug 18, 1987
HOSPIRA INC	EQ 40MG BASE/VIAL	A040793	001	Nov 25, 2008
	EQ 125MG BASE/VIAL	A040827	001	Nov 25, 2008

METHYLPREDNISOLONE

ELKINS SINN	EQ 125MG BASE/VIAL	A086906	002	
	EQ 500MG BASE/VIAL	A086906	003	
	EQ 1GM BASE/VIAL	A086906	004	
ORGANON USA INC	EQ 500MG BASE/VIAL	A087535	001	Jun 25, 1982
	EQ 1GM BASE/VIAL	A087535	002	Jun 25, 1982

METHYLPREDNISOLONE SODIUM SUCCINATE

ABRAXIS PHARM	EQ 40MG BASE/VIAL	A088676	001	Jun 08, 1984	
	EQ 40MG BASE/VIAL	A089143	001	Mar 28, 1986	
	EQ 125MG BASE/VIAL	A088677	001	Jun 08, 1984	
	EQ 125MG BASE/VIAL	A089144	001	Mar 28, 1986	
	EQ 500MG BASE/VIAL	A088678	001	Jun 08, 1984	
	EQ 500MG BASE/VIAL	A089186	001	Mar 28, 1986	
	EQ 500MG BASE/VIAL	A089187	001	Mar 28, 1986	
	EQ 1GM BASE/VIAL	A088679	001	Jun 08, 1984	
	EQ 1GM BASE/VIAL	A089188	001	Mar 28, 1986	
	EQ 1GM BASE/VIAL	A089189	001	Mar 28, 1986	
	BEDFORD LABS	EQ 40MG BASE/VIAL	A040662	001	Feb 21, 2007
		EQ 125MG BASE/VIAL	A040641	002	Feb 21, 2007
		EQ 500MG BASE/VIAL	A040641	003	Feb 21, 2007
EQ 500MG BASE/VIAL		A040709	001	Feb 21, 2007	
EQ 1GM BASE/VIAL		A040641	004	Feb 21, 2007	
ELKINS SINN	EQ 1GM BASE/VIAL	A040709	002	Feb 21, 2007	
	EQ 40MG BASE/VIAL	A086906	001		
EUGIA PHARMA	EQ 40MG BASE/VIAL	A207667	001	Dec 15, 2015	
	EQ 125MG BASE/VIAL	A207667	002	Dec 15, 2015	
	EQ 500MG BASE/VIAL	A207667	003	Dec 15, 2015	
	EQ 2GM BASE/VIAL	A207667	004	Dec 15, 2015	
INTL MEDICATION	EQ 40MG BASE/VIAL	A087812	001	Feb 09, 1983	
	EQ 125MG BASE/VIAL	A087813	001	Feb 09, 1983	
	EQ 500MG BASE/VIAL	A087851	001	Feb 09, 1983	
TEVA PARENTERAL	EQ 1GM BASE/VIAL	A087852	001	Feb 09, 1983	
	EQ 125MG BASE/VIAL	A081266	001	Nov 30, 1992	
	EQ 500MG BASE/VIAL	A081267	001	Nov 30, 1992	
WATSON LABS	EQ 1GM BASE/VIAL	A081268	001	Nov 30, 1992	
	EQ 40MG BASE/VIAL	A086953	001	Jul 22, 1982	
	EQ 125MG BASE/VIAL	A087030	001	Jul 22, 1982	
	EQ 500MG BASE/VIAL	A088523	001	Jul 24, 1984	
	EQ 1GM BASE/VIAL	A088524	001	Jul 24, 1984	

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

METHYLPREDNISOLONE; NEOMYCIN SULFATE

OINTMENT;OPHTHALMIC

NEO-MEDROL

PHARMACIA AND UPJOHN 0.1%;EQ 3.5MG BASE/GM

A060645 001

METHYLTESTOSTERONE

CAPSULE;ORAL

METHYLTESTOSTERONE

HEATHER 10MG

A084967 001

TESTRED

+ BAUSCH 10MG

A083976 001

VIRILON

CHARTWELL 10MG

A087750 001 Nov 24, 1982

TABLET;BUCCAL

ANDROID 5

VALEANT PHARM INTL 5MG

A087222 001

ORETON

SCHERING 10MG

A080281 001

TABLET;BUCCAL, SUBLINGUAL

METANDREN

NOVARTIS 5MG

N003240 004

10MG

N003240 001

10MG

N003240 005

25MG

N003240 003

METHYLTESTOSTERONE

IMPAX LABS 10MG

A084287 001

LILLY 10MG

A080256 001

25MG

A080256 002

PUREPAC PHARM 10MG

A080308 001

10MG

A080475 001

10MG

A080475 002

25MG

A080475 003

PVT FORM 5MG

A083836 001

TABLICAPS 10MG

A085125 001

USL PHARMA 10MG

A080271 001

TABLET;ORAL

ANDROID 10

VALEANT PHARMS NORTH 10MG

A086450 001

METHYLTESTOSTERONE

IMPAX LABS 25MG

A084310 001

INWOOD LABS 10MG

A080839 001

25MG

A080973 001

KV PHARM 10MG

A084312 001

LANNETT 10MG

A087092 001 Nov 05, 1982

25MG

A087111 001 Jan 27, 1983

PARKE DAVIS 10MG

A084244 001

25MG

A084241 001

PUREPAC PHARM 10MG

A080309 001

25MG

A080310 001

PVT FORM 5MG

A080214 001

10MG

A080214 002

25MG

A080214 003

TABLICAPS 10MG

A080313 001

25MG

A085270 001

WATSON LABS 10MG

A080933 001

25MG

A080931 001

WEST WARD 10MG

A084331 001

25MG

A084331 002

25MG

A084642 001

ORETON METHYL

SCHERING 10MG

N003158 001

25MG

N003158 002

METHYPRYLON

CAPSULE;ORAL

NOLUDAR

ROCHE 300MG

N009660 008

ELIXIR;ORAL

NOLUDAR

ROCHE 50MG/5ML

N009660 007

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

METHYPRYLON

TABLET; ORAL

NOLUDAR

ROCHE	50MG	N009660	002
	200MG	N009660	004

METHYSERGIDE MALEATE

TABLET; ORAL

SANSERT

NOVARTIS	2MG	N012516	001
----------	-----	---------	-----

METIPRANOLOL HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

METIPRANOLOL

SANDOZ	0.3%	A075720	001	Aug 06, 2001
--------	------	---------	-----	--------------

OPTIPRANOLOL

+ BAUSCH AND LOMB	0.3% **	N019907	001	Dec 29, 1989
-------------------	---------	---------	-----	--------------

METOCLOPRAMIDE HYDROCHLORIDE

CONCENTRATE; ORAL

METOCLOPRAMIDE INTENSOL

ROXANE	EQ 10MG BASE/ML	A072995	001	Jan 30, 1992
--------	-----------------	---------	-----	--------------

INJECTABLE; INJECTION

METOCLOPRAMIDE HYDROCHLORIDE

BEDFORD	EQ 5MG BASE/ML	A072155	001	Mar 30, 1992
---------	----------------	---------	-----	--------------

	EQ 5MG BASE/ML	A072244	001	Mar 30, 1992
--	----------------	---------	-----	--------------

	EQ 5MG BASE/ML	A072247	001	May 18, 1992
--	----------------	---------	-----	--------------

HOSPIRA	EQ 5MG BASE/ML	A070505	001	Jun 23, 1989
---------	----------------	---------	-----	--------------

	EQ 5MG BASE/ML	A070506	001	Jun 22, 1989
--	----------------	---------	-----	--------------

	EQ 5MG BASE/ML	A070847	001	Nov 07, 1988
--	----------------	---------	-----	--------------

	EQ 5MG BASE/ML	A071291	001	Mar 03, 1989
--	----------------	---------	-----	--------------

	EQ 5MG BASE/ML	A071990	001	Jan 18, 1989
--	----------------	---------	-----	--------------

	EQ 5MG BASE/ML	A073117	001	Jan 17, 1991
--	----------------	---------	-----	--------------

	EQ 5MG BASE/ML	A074147	001	Aug 02, 1996
--	----------------	---------	-----	--------------

LYPHOMED	EQ 10MG BASE/2ML	A070293	001	Jan 24, 1986
----------	------------------	---------	-----	--------------

NORBROOK	EQ 10MG BASE/2ML	A070892	001	Aug 26, 1988
----------	------------------	---------	-----	--------------

SMITH AND NEPHEW	EQ 5MG BASE/ML	A070623	001	Mar 02, 1987
------------------	----------------	---------	-----	--------------

	EQ 10MG BASE/2ML	A070622	001	Mar 02, 1987
--	------------------	---------	-----	--------------

REGLAN

+ HIKMA	EQ 5MG BASE/ML **	N017862	001
---------	-------------------	---------	-----

	EQ 10MG BASE/ML **	N017862	004	May 28, 1987
--	--------------------	---------	-----	--------------

SOLUTION; ORAL

METOCLOPRAMIDE HYDROCHLORIDE

ACTAVIS MID ATLANTIC	EQ 5MG BASE/5ML	A071340	001	Aug 18, 1988
----------------------	-----------------	---------	-----	--------------

CHARTWELL MOLECULAR	EQ 5MG BASE/5ML	A073680	001	Oct 27, 1992
---------------------	-----------------	---------	-----	--------------

MORTON GROVE	EQ 5MG BASE/5ML	A070949	001	Mar 06, 1987
--------------	-----------------	---------	-----	--------------

PACO	EQ 5MG BASE/5ML	A071665	001	Dec 05, 1988
------	-----------------	---------	-----	--------------

PHARMOBEDIANT CNSLTG	EQ 5MG BASE/5ML	A074703	001	Oct 31, 1997
----------------------	-----------------	---------	-----	--------------

ROXANE	EQ 5MG BASE/5ML	A072038	001	Dec 05, 1988
--------	-----------------	---------	-----	--------------

TEVA	EQ 5MG BASE/5ML	A070819	001	Jul 10, 1987
------	-----------------	---------	-----	--------------

	EQ 5MG BASE/5ML	A071315	001	Jun 30, 1993
--	-----------------	---------	-----	--------------

VISTAPHARM	EQ 5MG BASE/5ML	A075051	001	Jan 26, 2001
------------	-----------------	---------	-----	--------------

REGLAN

+ ROBINS AH	EQ 5MG BASE/5ML **	N018821	001	Mar 25, 1983
-------------	--------------------	---------	-----	--------------

TABLET; ORAL

CLOPRA

QUANTUM PHARMICS	EQ 5MG BASE	A072384	001	Jun 02, 1988
------------------	-------------	---------	-----	--------------

	EQ 10MG BASE	A070294	001	Jul 29, 1985
--	--------------	---------	-----	--------------

CLOPRA-"YELLOW"

QUANTUM PHARMICS	EQ 10MG BASE	A070632	001	Oct 28, 1985
------------------	--------------	---------	-----	--------------

MAXOLON

KING PHARMS	EQ 10MG BASE	A070106	001	Mar 04, 1986
-------------	--------------	---------	-----	--------------

METOCLOPRAMIDE HYDROCHLORIDE

CHARTWELL RX	EQ 5MG BASE	A074478	001	Oct 05, 1995
--------------	-------------	---------	-----	--------------

	EQ 10MG BASE	A074478	002	Oct 05, 1995
--	--------------	---------	-----	--------------

CLONMEL	EQ 10MG BASE	A072639	001	May 09, 1991
---------	--------------	---------	-----	--------------

HALSEY	EQ 10MG BASE	A070906	001	Oct 28, 1986
--------	--------------	---------	-----	--------------

INTERPHARM	EQ 10MG BASE	A071213	001	Sep 24, 1986
------------	--------------	---------	-----	--------------

MUTUAL PHARM	EQ 10MG BASE	A070660	001	Feb 10, 1987
--------------	--------------	---------	-----	--------------

NORTHSTAR HLTHCARE	EQ 5MG BASE	A078374	001	Nov 30, 2007
--------------------	-------------	---------	-----	--------------

	EQ 10MG BASE	A078374	002	Nov 30, 2007
--	--------------	---------	-----	--------------

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

METOCLOPRAMIDE HYDROCHLORIDE

TABLET; ORAL

METOCLOPRAMIDE HYDROCHLORIDE

PAR PHARM	EQ 10MG BASE	A070342	001	Mar 25, 1986
SANDOZ	EQ 5MG BASE	A072436	001	Jun 22, 1989
	EQ 10MG BASE	A070850	001	Feb 03, 1987
SCHERING	EQ 10MG BASE	A070598	001	Feb 02, 1987
SUN PHARM INDUSTRIES	EQ 5MG BASE	A071536	002	Jan 16, 1997
	EQ 10MG BASE	A071536	001	Apr 28, 1993
SUPERPHARM	EQ 10MG BASE	A070926	001	Jun 26, 1987
USL PHARMA	EQ 10MG BASE	A070339	001	Jul 29, 1985
WATSON LABS	EQ 10MG BASE	A070363	001	Mar 02, 1987
	EQ 10MG BASE	A070453	001	Jun 06, 1986
	EQ 10MG BASE	A070511	001	Jan 22, 1986
	EQ 10MG BASE	A070645	001	May 11, 1987

TABLET, ORALLY DISINTEGRATING; ORAL

METOZOLV ODT

+ SALIX PHARMS	EQ 5MG BASE **	N022246	001	Sep 04, 2009
+	EQ 10MG BASE **	N022246	002	Sep 04, 2009

REGLAN ODT

MEDA PHARMS	EQ 5MG BASE	N021793	001	Jun 10, 2005
	EQ 10MG BASE	N021793	002	Jun 10, 2005

METOCURINE IODIDE

INJECTABLE; INJECTION

METUBINE IODIDE

LILLY	2MG/ML	N006632	003	
-------	--------	---------	-----	--

METOLAZONE

TABLET; ORAL

DIULO

GD SEARLE LLC	2.5MG	N018535	001	
	5MG	N018535	002	
	10MG	N018535	003	

METOLAZONE

ANI PHARMS	2.5MG	A075543	001	Jan 06, 2004
	5MG	A075543	002	Mar 01, 2004
	10MG	A075543	003	Dec 24, 2003
ROXANE	10MG	A076482	002	Apr 29, 2004
WATSON LABS	10MG	A076891	001	Jul 21, 2004

MYKROX

CHARTWELL MOLECULAR	0.5MG	N019532	001	Oct 30, 1987
---------------------	-------	---------	-----	--------------

ZAROXOLYN

+ I3 PHARMS	2.5MG **	N017386	001	
+	5MG **	N017386	002	
+	10MG **	N017386	003	

METOPROLOL FUMARATE

TABLET, EXTENDED RELEASE; ORAL

LOPRESSOR

NOVARTIS	EQ 100MG TARTRATE	N019786	001	Dec 27, 1989
	EQ 200MG TARTRATE	N019786	002	Dec 27, 1989
	EQ 300MG TARTRATE	N019786	003	Dec 27, 1989
	EQ 400MG TARTRATE	N019786	004	Dec 27, 1989

METOPROLOL SUCCINATE

TABLET, EXTENDED RELEASE; ORAL

METOPROLOL SUCCINATE

ACCORD HLTHCARE	EQ 25MG TARTRATE	A215637	001	Oct 18, 2022
	EQ 50MG TARTRATE	A215637	002	Oct 18, 2022
	EQ 100MG TARTRATE	A215637	003	Oct 18, 2022
	EQ 200MG TARTRATE	A215637	004	Oct 18, 2022
ACTAVIS LABS FL INC	EQ 25MG TARTRATE	A076862	002	Aug 03, 2009
	EQ 100MG TARTRATE	A077298	001	Apr 15, 2010
	EQ 200MG TARTRATE	A077298	002	Apr 15, 2010
GRANULES	EQ 25MG TARTRATE	A216509	001	Aug 07, 2023
	EQ 50MG TARTRATE	A216509	002	Aug 07, 2023
	EQ 100MG TARTRATE	A216509	003	Aug 07, 2023
	EQ 200MG TARTRATE	A216509	004	Aug 07, 2023
LUPIN	EQ 25MG TARTRATE	A209272	001	Aug 15, 2023
	EQ 50MG TARTRATE	A209272	002	Aug 15, 2023

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

METOPROLOL SUCCINATETABLET, EXTENDED RELEASE;ORAL
METOPROLOL SUCCINATE

	EQ 100MG TARTRATE	A209272 003	Aug 15, 2023
	EQ 200MG TARTRATE	A209272 004	Aug 15, 2023
NESHER PHARMS	EQ 25MG TARTRATE	A077779 001	Mar 20, 2008
	EQ 50MG TARTRATE	A077176 001	May 14, 2008
	EQ 100MG TARTRATE	A076640 002	May 18, 2007
	EQ 200MG TARTRATE	A076640 001	May 18, 2007
PRINSTON INC	EQ 100MG TARTRATE	A210597 001	Jan 04, 2022
	EQ 200MG TARTRATE	A210597 002	Jan 04, 2022
SANDOZ	EQ 25MG TARTRATE	A076969 001	Jul 31, 2006
	EQ 50MG TARTRATE	A076969 002	May 18, 2007
	EQ 100MG TARTRATE	A076969 003	Mar 20, 2008
	EQ 200MG TARTRATE	A076969 004	Mar 20, 2008

METOPROLOL TARTRATE

INJECTABLE;INJECTION

LOPRESSOR

+ NOVARTIS

1MG/ML **

N018704 001 Mar 30, 1984

METOPROLOL TARTRATE

AM REGENT

1MG/ML

A090386 001 Sep 30, 2009

HOSPIRA

1MG/ML

A074133 001 Dec 21, 1993

1MG/ML

A075160 001 Jul 06, 1998

LUITPOLD

1MG/ML

A091307 001 Dec 29, 2010

STERISCIENCE SPECLTS

1MG/ML

A090317 001 Apr 19, 2010

WATSON LABS

1MG/ML

A074032 001 Dec 21, 1993

TABLET;ORAL

METOPROLOL TARTRATE

APOTHECON

50MG

A074258 001 Jan 27, 1994

100MG

A074258 002 Jan 27, 1994

CHARTWELL RX

50MG

A073288 001 Mar 25, 1994

100MG

A073289 001 Mar 25, 1994

HERITAGE PHARMA

50MG

A074141 001 Jan 31, 1995

100MG

A074141 002 Jan 31, 1995

NORVIUM BIOSCIENCE

50MG

A073666 001 Dec 21, 1993

100MG

A073666 002 Dec 21, 1993

PUREPAC PHARM

50MG

A074380 001 Jul 29, 1994

100MG

A074380 002 Jul 29, 1994

SUN PHARM INDUSTRIES

25MG

A073654 002 Jul 15, 2009

50MG

A073654 003 Dec 21, 1993

100MG

A073654 001 Dec 21, 1993

TEVA

50MG

A074143 001 Sep 30, 1994

100MG

A074143 002 Sep 30, 1994

TEVA PHARMS

50MG

A074333 001 Jan 27, 1994

100MG

A074333 002 Jan 27, 1994

WATSON LABS

50MG

A074217 001 May 27, 1994

100MG

A074217 002 May 27, 1994

ZYDUS PHARMS

25MG

A212402 001 Apr 19, 2023

50MG

A212402 002 Apr 19, 2023

100MG

A212402 003 Apr 19, 2023

METRIZAMIDE

INJECTABLE;INJECTION

AMIPAQUE

GE HEALTHCARE

2.5GM/VIAL

N017982 003 Sep 12, 1983

3.75GM/VIAL

N017982 001

6.75GM/VIAL

N017982 002

13.5GM/VIAL

N017982 004 Sep 12, 1983

METRONIDAZOLE

CAPSULE;ORAL

METRONIDAZOLE

ABLE

375MG

A076505 001 Nov 13, 2003

GEL;TOPICAL

METROGEL

+ GALDERMA LABS LP

0.75%

N019737 001 Nov 22, 1988

METRONIDAZOLE

CHARTWELL RX

1%

A090903 001 Jul 22, 2011

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

METRONIDAZOLE

GEL;VAGINAL

METRONIDAZOLE

ENCUBE 0.75% A077264 001 Oct 31, 2006

INJECTABLE;INJECTION

FLAGYL I.V. RTU IN PLASTIC CONTAINER

PFIZER 500MG/100ML N018353 002

METRO I.V.

B BRAUN 500MG/100ML N018674 001 Aug 31, 1982

METRONIDAZOLE

ABBOTT 500MG/100ML N018889 001 Nov 18, 1983

ABRAXIS PHARM 500MG/100ML A070071 001 Dec 03, 1984

HIKMA 500MG/100ML N018907 001 Mar 30, 1984

INTL MEDICATION 500MG/100ML A070004 001 May 08, 1985

WATSON LABS 500MG/100ML A070042 001 Dec 20, 1984

500MG/100ML A070170 001 Apr 01, 1986

METRONIDAZOLE IN PLASTIC CONTAINER

RISING 500MG/100ML A205531 001 May 09, 2017

TABLET;ORAL

FLAGYL

+ PFIZER 250MG ** N012623 001

+ 500MG ** N012623 003

METROMIDOL

LABS AF 250MG A074523 001 Oct 24, 1996

500MG A074523 002 Oct 24, 1996

METRONIDAZOLE

ABLE 250MG A076519 001 Jun 27, 2003

500MG A076519 002 Jun 27, 2003

CHARTWELL MOLECULES 250MG N018845 001 Aug 18, 1983

500MG N018930 001 Aug 18, 1983

FLAMINGO PHARMS 250MG A207309 001 May 16, 2016

500MG A207309 002 May 16, 2016

FOSUN PHARMA 250MG N018620 001 Mar 04, 1982

250MG N018740 001 Oct 22, 1982

500MG N018620 002 Jun 02, 1983

500MG N018740 002 Oct 22, 1982

HALSEY 250MG A070021 001 Apr 02, 1985

500MG A070593 001 Feb 27, 1986

IVAX SUB TEVA PHARMS 250MG N018517 001

500MG N018517 002 May 05, 1982

LNK 250MG N019029 001 Apr 10, 1984

MUTUAL PHARM 250MG N018818 001 Feb 16, 1983

500MG N018818 002 Feb 16, 1983

SUPERPHARM 250MG A070008 001 Dec 11, 1984

500MG A070009 001 Dec 11, 1984

WATSON LABS 250MG N018599 001 Sep 17, 1982

250MG N018764 001 Sep 17, 1982

500MG N018599 002 Feb 13, 1984

500MG N018764 002 Dec 20, 1982

PROTOSTAT

ORTHO MCNEIL PHARM 250MG N018871 001 Mar 02, 1983

500MG N018871 002 Mar 02, 1983

SATRIC

SAVAGE LABS 250MG A070029 001 Mar 19, 1985

500MG A070731 001 Jun 08, 1987

TABLET, EXTENDED RELEASE;ORAL

FLAGYL ER

+ PFIZER 750MG N020868 001 Nov 26, 1997

METRONIDAZOLE

ABLE 750MG A076462 001 Jun 25, 2003

ALEMBIC 750MG A090222 001 May 05, 2010

METRONIDAZOLE HYDROCHLORIDE

INJECTABLE;INJECTION

FLAGYL I.V.

PFIZER EQ 500MG BASE/VIAL ** N018353 001

METRONIDAZOLE HYDROCHLORIDE

ABRAXIS PHARM EQ 500MG BASE/VIAL A070295 001 Oct 15, 1985

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

METYRAPONETABLET; ORAL
METOPIRONE

HRA PHARMA 250MG N012911 001

MEXILETINE HYDROCHLORIDE

CAPSULE; ORAL

MEXILETINE HYDROCHLORIDE

WATSON LABS	150MG	A074711 001	Feb 26, 1997
	150MG	A074865 001	Apr 13, 1998
	200MG	A074711 002	Feb 26, 1997
	200MG	A074865 002	Apr 13, 1998
	250MG	A074711 003	Feb 26, 1997
	250MG	A074865 003	Apr 13, 1998

MEXITIL

+	BOEHRINGER INGELHEIM	150MG **	N018873 002	Dec 30, 1985
+		200MG **	N018873 003	Dec 30, 1985
+		250MG **	N018873 004	Dec 30, 1985

MEZLOCILLIN SODIUM MONOHYDRATE

INJECTABLE; INJECTION

MEZLIN

BAYER PHARMS	EQ 1GM BASE/VIAL	A062333 001	
	EQ 1GM BASE/VIAL	A062372 005	Jan 13, 1983
	EQ 1GM BASE/VIAL	N050549 001	
	EQ 2GM BASE/VIAL	A062333 002	
	EQ 2GM BASE/VIAL	A062372 001	May 13, 1982
	EQ 2GM BASE/VIAL	N050549 002	
	EQ 3GM BASE/VIAL	A062333 003	
	EQ 3GM BASE/VIAL	A062372 002	May 13, 1982
	EQ 3GM BASE/VIAL	A062697 001	Jan 22, 1987
	EQ 3GM BASE/VIAL	N050549 003	
	EQ 4GM BASE/VIAL	A062333 004	
	EQ 4GM BASE/VIAL	A062372 003	May 13, 1982
	EQ 4GM BASE/VIAL	A062697 002	Jan 22, 1987
	EQ 4GM BASE/VIAL	N050549 004	
	EQ 20GM BASE/VIAL	A062372 004	Mar 02, 1988
	EQ 20GM BASE/VIAL	N050549 005	Mar 02, 1988

MICAFUNGIN SODIUM

INJECTABLE; INTRAVENOUS

MICAFUNGIN SODIUM

APOTEX	EQ 50MG BASE/VIAL	A208366 001	Nov 05, 2020
	EQ 100MG BASE/VIAL	A208366 002	Nov 05, 2020

POWDER; INTRAVENOUS

MICAFUNGIN

+	TEVA PHARMS USA INC	EQ 50MG BASE/VIAL	N212125 001	Jul 30, 2021
+		EQ 100MG BASE/VIAL	N212125 002	Jul 30, 2021

MICONAZOLE

INJECTABLE; INJECTION

MONISTAT

JANSSEN PHARMA 10MG/ML N018040 001

MICONAZOLE NITRATE

CREAM; TOPICAL

MONISTAT-DERM

INSIGHT PHARMS 2% N017494 001

CREAM; VAGINAL

MICONAZOLE NITRATE

COSETTE	2%	A074366 001	Feb 22, 1996
TEVA	2%	A074136 001	Jan 04, 1995
TEVA PHARMS	2%	A074030 001	Oct 30, 1992

CREAM, SUPPOSITORY; TOPICAL, VAGINAL

M-ZOLE 7 DUAL PACK

ACTAVIS MID ATLANTIC 2%, 100MG A074586 001 Jul 17, 1997

MICONAZOLE 7 COMBINATION PACK

COSETTE 2%, 100MG A076585 001 Mar 26, 2004

LOTION; TOPICAL

MONISTAT-DERM

INSIGHT PHARMS 2% N017739 001

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

MICONAZOLE NITRATE

SUPPOSITORY; VAGINAL

MICONAZOLE NITRATE

COSETTE	100MG	A074414	001	Apr 30, 1997
PERRIGO	100MG	A074395	001	Mar 20, 1997

TAMPON; VAGINAL

MONISTAT 5

PERSONAL PRODS	100MG	N018592	001	Oct 27, 1989
----------------	-------	---------	-----	--------------

MIDAZOLAM

SOLUTION; INTRAVENOUS

MIDAZOLAM IN 0.8% SODIUM CHLORIDE

+ EXELA PHARMA	50MG/50ML (1MG/ML) **	N215868	001	Jul 20, 2022
+	100MG/100ML (1MG/ML)	N215868	002	Jul 20, 2022

MIDAZOLAM HYDROCHLORIDE

INJECTABLE; INJECTION

MIDAZOLAM HYDROCHLORIDE

APOTHECON	EQ 1MG BASE/ML	A075620	001	Nov 01, 2000
	EQ 5MG BASE/ML	A075620	002	Nov 01, 2000
	EQ 5MG BASE/ML	A075641	001	Oct 19, 2000
BAXTER HLTHCARE CORP	EQ 1MG BASE/ML	A075637	001	Oct 31, 2000
	EQ 5MG BASE/ML	A075637	002	Oct 31, 2000
BEDFORD	EQ 5MG BASE/ML	A075249	001	Jun 23, 2000
BEN VENUE	EQ 5MG BASE/ML	A075455	001	Jun 20, 2000
ENDO OPERATIONS	EQ 1MG BASE/ML	A078141	001	May 30, 2008
	EQ 1MG BASE/ML	A078511	001	Nov 10, 2008
	EQ 5MG BASE/ML	A078141	002	May 30, 2008
	EQ 5MG BASE/ML	A078511	002	Nov 10, 2008
FRESENIUS KABI USA	EQ 5MG BASE/ML	A208878	001	Mar 28, 2017
HOSPIRA	EQ 1MG BASE/ML	A075396	001	Jun 20, 2000
	EQ 1MG BASE/ML	A075856	001	Jun 13, 2002
	EQ 5MG BASE/ML	A075396	002	Jun 20, 2000
	EQ 5MG BASE/ML	A075484	001	Jun 20, 2000
	EQ 5MG BASE/ML	A075856	002	Jun 13, 2002
HOSPIRA INC	EQ 1MG BASE/ML	A075409	002	Jun 20, 2000
	EQ 5MG BASE/ML	A075409	001	Jun 20, 2000
IGI LABS INC	EQ 5MG BASE/ML	A075263	001	Jun 26, 2000
INTL MEDICATED	EQ 1MG BASE/ML	A076144	001	Jan 26, 2005
	EQ 5MG BASE/ML	A076144	002	Jan 26, 2005
INTL MEDICATION	EQ 1MG BASE/ML	A076020	001	Jul 16, 2004
	EQ 5MG BASE/ML	A076020	002	Jul 16, 2004
RISING	EQ 5MG BASE/ML	A075481	001	Jun 30, 2000
VERSED				
+ HLR	EQ 1MG BASE/ML **	N018654	002	May 26, 1987
+	EQ 5MG BASE/ML **	N018654	001	Dec 20, 1985

SYRUP; ORAL

MIDAZOLAM HYDROCHLORIDE

PAI HOLDINGS	EQ 2MG BASE/ML	A075958	001	Sep 04, 2003
PHARM ASSOC	EQ 2MG BASE/ML	A077115	001	Sep 09, 2005
SUN PHARM INDS LTD	EQ 2MG BASE/ML	A076058	001	Mar 15, 2002

VERSED

+ ROCHE	EQ 2MG BASE/ML **	N020942	001	Oct 15, 1998
---------	-------------------	---------	-----	--------------

MIDODRINE HYDROCHLORIDE

TABLET; ORAL

MIDODRINE HYDROCHLORIDE

CHARTWELL RX	2.5MG	A076514	001	Sep 11, 2003
	5MG	A076514	002	Sep 11, 2003
	10MG	A076514	003	Jul 02, 2004

PROAMATINE

+ TAKEDA PHARMS USA	2.5MG **	N019815	001	Sep 06, 1996
+	5MG **	N019815	002	Sep 06, 1996
+	10MG **	N019815	003	Mar 20, 2002

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

MIDOSTAURIN

CAPSULE; ORAL

MIDOSTAURIN

TEVA PHARMS 25MG A216076 001 Apr 29, 2024

MIGLUSTAT

CAPSULE; ORAL

MIGLUSTAT

BRECKENRIDGE 100MG A209325 001 Feb 03, 2022

MILNACIPRAN HYDROCHLORIDE

TABLET; ORAL

MILNACIPRAN HYDROCHLORIDE

AMNEAL PHARMS 12.5MG A205081 001 Apr 22, 2016
 25MG A205081 002 Apr 22, 2016
 50MG A205081 003 Apr 22, 2016
 100MG A205081 004 Apr 22, 2016
 FIRST TIME US 12.5MG A205071 001 Jan 27, 2016
 25MG A205071 002 Jan 27, 2016
 50MG A205071 003 Jan 27, 2016
 100MG A205071 004 Jan 27, 2016

MILRINONE LACTATE

INJECTABLE; INJECTION

MILRINONE LACTATE

BAXTER HLTHCARE CORP EQ 1MG BASE/ML A076427 001 Sep 21, 2004
 GLAND PHARMA LTD EQ 1MG BASE/ML A077190 001 Oct 31, 2006
 HIKMA EQ 1MG BASE/ML A075852 001 May 28, 2002
 HOSPIRA EQ 1MG BASE/ML A075830 001 May 28, 2002
 EQ 1MG BASE/ML A075884 001 May 28, 2002
 INTL MEDICATED EQ 1MG BASE/ML A076013 001 Aug 02, 2002
 MYLAN INSTITUTIONAL EQ 1MG BASE/ML A076428 001 Jun 16, 2003

MILRINONE LACTATE IN DEXTROSE 5%

WOODWARD EQ 40MG BASE/200ML (EQ 0.2MG BASE/ML) A077151 002 Jul 20, 2005

MILRINONE LACTATE IN DEXTROSE 5% IN PLASTIC CONTAINER

B BRAUN EQ 20MG BASE/100ML (EQ 0.2MG BASE/ML) A076414 001 Aug 18, 2004
 BAXTER HLTHCARE EQ 20MG BASE/100ML (EQ 0.2MG BASE/ML) A076259 001 Aug 08, 2002
 HIKMA EQ 20MG BASE/100ML (EQ 0.2MG BASE/ML) A075510 001 May 28, 2002
 HOSPIRA EQ 20MG BASE/100ML (EQ 0.2MG BASE/ML) A075885 001 May 28, 2002
 EQ 40MG BASE/200ML (EQ 0.2MG BASE/ML) A075885 002 May 28, 2002
 WOODWARD EQ 20MG BASE/100ML (EQ 0.2MG BASE/ML) A077151 001 Jul 20, 2005

PRIMACOR

+ SANOFI AVENTIS US EQ 1MG BASE/ML ** N019436 001 Dec 31, 1987

PRIMACOR IN DEXTROSE 5% IN PLASTIC CONTAINER

+ SANOFI AVENTIS US EQ 10MG BASE/100ML ** N020343 001 Aug 09, 1994

+ EQ 15MG BASE/100ML ** N020343 002 Aug 09, 1994

+ EQ 20MG BASE/100ML (EQ 0.2MG BASE/ML) N020343 003 Aug 09, 1994

**

+ EQ 40MG BASE/200ML (EQ 0.2MG BASE/ML) N020343 004 Aug 09, 1994

**

MINOCYCLINE HYDROCHLORIDE

CAPSULE; ORAL

DYNACIN

ALVOGEN EQ 50MG BASE A063067 003 Aug 14, 1990
 EQ 75MG BASE A063067 002 Sep 15, 1999
 EQ 100MG BASE A063067 001 Jul 31, 1990

MINOCIN

+ BAUSCH EQ 75MG BASE ** N050649 003 Feb 12, 2001

TRIAx PHARMS EQ 50MG BASE N050315 002

EQ 100MG BASE N050315 001

CAPSULE, EXTENDED RELEASE; ORAL

XIMINO

JOURNEY EQ 45MG BASE N201922 001 Jul 11, 2012
 EQ 67.5MG BASE N201922 002 Jul 11, 2012
 EQ 90MG BASE N201922 003 Jul 11, 2012
 EQ 112.5MG BASE N201922 004 Jul 11, 2012
 EQ 135MG BASE N201922 005 Jul 11, 2012

INJECTABLE; INJECTION

MINOCIN

LEDERLE EQ 100MG BASE/VIAL A062139 001

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

MINOCYCLINE HYDROCHLORIDE

SUSPENSION;ORAL

MINOCIN

BAUSCH EQ 50MG BASE/5ML N050445 001

TABLET;ORAL

MINOCYCLINE HYDROCHLORIDE

SUN PHARM INDUSTRIES EQ 50MG BASE A090217 001 Jan 29, 2016

EQ 75MG BASE A090217 002 Jan 29, 2016

EQ 100MG BASE A090217 003 Jan 29, 2016

+ TRIAX PHARMS EQ 50MG BASE ** N050451 003 Aug 10, 1982

+ EQ 100MG BASE ** N050451 002 Aug 10, 1982

TABLET, EXTENDED RELEASE;ORAL

MINOCYCLINE HYDROCHLORIDE

BARR LABS INC EQ 45MG BASE A065485 001 Mar 17, 2009

EQ 65MG BASE A065485 004 May 18, 2012

EQ 80MG BASE A065485 007 Apr 26, 2017

EQ 90MG BASE A065485 002 Mar 17, 2009

EQ 105MG BASE A065485 008 Apr 26, 2017

EQ 115MG BASE A065485 005 May 18, 2012

EQ 135MG BASE A065485 003 Mar 17, 2009

IMPAX LABS INC EQ 45MG BASE A090024 001 Feb 03, 2009

EQ 90MG BASE A090024 002 Feb 03, 2009

EQ 135MG BASE A090024 003 Feb 03, 2009

NORVIUM BIOSCIENCE EQ 45MG BASE A090911 001 Jul 20, 2010

EQ 90MG BASE A090911 002 Jul 20, 2010

EQ 135MG BASE A090911 003 Jul 20, 2010

RISING EQ 55MG BASE A203443 001 Aug 21, 2019

EQ 65MG BASE A201467 001 Jul 30, 2019

EQ 80MG BASE A203443 002 Aug 22, 2014

EQ 105MG BASE A203443 003 Aug 22, 2014

EQ 115MG BASE A201467 002 Jul 30, 2019

SUN PHARM INDS LTD EQ 45MG BASE A091118 001 Sep 25, 2014

EQ 65MG BASE A091118 003 Dec 03, 2019

EQ 80MG BASE A091118 004 Sep 25, 2014

EQ 90MG BASE A091118 005 Sep 25, 2014

EQ 105MG BASE A091118 006 Sep 25, 2014

EQ 115MG BASE A091118 007 Dec 03, 2019

EQ 135MG BASE A091118 008 Sep 25, 2014

ZYDUS PHARMS EQ 45MG BASE A203553 001 Nov 16, 2017

EQ 55MG BASE A203553 002 Jun 16, 2023

EQ 65MG BASE A203553 003 Jun 16, 2023

EQ 80MG BASE A203553 004 Nov 16, 2017

EQ 90MG BASE A203553 005 Nov 16, 2017

EQ 105MG BASE A203553 006 Nov 16, 2017

EQ 115MG BASE A203553 007 Jun 16, 2023

EQ 135MG BASE A203553 008 Nov 16, 2017

MINOLIRA

EPI HLTH EQ 105MG BASE N209269 001 May 08, 2017

EQ 135MG BASE N209269 002 May 08, 2017

SOLODYN

+ BAUSCH EQ 45MG BASE ** N050808 001 May 08, 2006

+ EQ 55MG BASE N050808 008 Aug 27, 2010

+ EQ 65MG BASE N050808 004 Jul 23, 2009

+ EQ 80MG BASE N050808 007 Aug 27, 2010

+ EQ 90MG BASE ** N050808 002 May 08, 2006

+ EQ 105MG BASE N050808 006 Aug 27, 2010

+ EQ 115MG BASE N050808 005 Jul 23, 2009

+ EQ 135MG BASE ** N050808 003 May 08, 2006

MINOXIDIL

SOLUTION;TOPICAL

MINOXIDIL (FOR MEN)

APOTEX INC 2% A074924 001 Apr 29, 1998

AUROBINDO PHARMA LTD 2% A074767 001 Feb 28, 1997

BAUSCH AND LOMB 2% A074643 001 Apr 09, 1996

COPLEY PHARM 2% A074500 001 May 23, 1996

HIKMA 2% A074731 001 Dec 24, 1996

SIGHT PHARMS 2% A074743 002 Oct 18, 1996

TEVA 2% A074589 001 Apr 05, 1996

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

MINOXIDIL

SOLUTION;TOPICAL

MINOXIDIL (FOR WOMEN)

APOTEX INC	2%	A074924	002	Apr 29, 1998
AUROBINDO PHARMA LTD	2%	A074767	002	Nov 25, 1997
HIKMA	2%	A074731	002	May 11, 2005
SIGHT PHARMS	2%	A074743	001	Oct 18, 1996

MINOXIDIL EXTRA STRENGTH (FOR MEN)

APOTEX INC	5%	A075839	001	Oct 01, 2001
AVACOR PRODS	5%	A075619	001	Nov 17, 2000
PERRIGO NEW YORK	5%	A075737	001	Mar 15, 2002

TABLET;ORAL

LONITEN

+ PFIZER	2.5MG **	N018154	001	
+	10MG **	N018154	003	

MINODYL

QUANTUM PHARMICS	2.5MG	A072153	001	Jul 13, 1988
	10MG	A071534	001	Mar 19, 1987

MINOXIDIL

ROYCE LABS	2.5MG	A071799	001	Nov 10, 1987
	10MG	A071796	001	Nov 10, 1987
USL PHARMA	2.5MG	A071537	001	Dec 16, 1988

MIPOMERSEN SODIUM

SOLUTION;SUBCUTANEOUS

KYNAMRO

+ KASTLE THERAPS LLC	200MG/ML (200MG/ML)	N203568	001	Jan 29, 2013
----------------------	---------------------	---------	-----	--------------

MIRABEGRON

TABLET, EXTENDED RELEASE;ORAL

MIRABEGRON

SAWAI USA	25MG	A209446	001	Dec 27, 2019
-----------	------	---------	-----	--------------

MIRTAZAPINE

TABLET;ORAL

MIRTAZAPINE

ACTAVIS ELIZABETH	15MG	A076241	001	Jun 25, 2003
	15MG	A076308	001	Jun 20, 2003
	30MG	A076241	002	Jun 25, 2003
	30MG	A076308	002	Jun 20, 2003
	45MG	A076241	003	Jun 25, 2003
	45MG	A076308	003	Jun 20, 2003
ACTAVIS LABS FL INC	15MG	A076336	001	Jun 20, 2003
	30MG	A076336	002	Jun 20, 2003
	45MG	A076336	003	Jun 20, 2003
IVAX SUB TEVA PHARMS	15MG	A076244	001	Dec 22, 2003
	30MG	A076244	002	Dec 22, 2003
	45MG	A076244	003	Dec 22, 2003
NORVIUM BIOSCIENCE	15MG	A076176	001	Jun 19, 2003
	30MG	A076176	002	Jun 19, 2003
	45MG	A076176	003	Jun 19, 2003
ROXANE	15MG	A076270	001	Jun 19, 2003
	30MG	A076270	002	Jun 19, 2003
	45MG	A076270	003	Jun 19, 2003
UPSHER SMITH LABS	15MG	A076189	001	Jun 19, 2003
	30MG	A076189	002	Jun 19, 2003
	45MG	A076189	003	Jun 19, 2003
WATSON LABS	15MG	A076312	001	Jun 19, 2003
	30MG	A076312	002	Jun 19, 2003
	45MG	A076312	003	Jun 19, 2003

REMERON

+ ORGANON	45MG **	N020415	003	Mar 17, 1997
-----------	---------	---------	-----	--------------

TABLET, ORALLY DISINTEGRATING;ORAL

MIRTAZAPINE

ACTAVIS ELIZABETH	15MG	A076689	001	Aug 31, 2005
	15MG	A077959	001	Feb 14, 2011
	30MG	A076689	002	Aug 31, 2005
	30MG	A077959	002	Feb 14, 2011
	45MG	A076689	003	Aug 31, 2005
	45MG	A077959	003	Feb 14, 2011
ACTAVIS LABS FL INC	15MG	A076307	001	Dec 17, 2003

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

MIRTAZAPINETABLET, ORALLY DISINTEGRATING;ORAL
MIRTAZAPINE

	30MG	A076307 002	Dec 17, 2003
	45MG	A076307 003	Feb 28, 2006
IMPAX LABS INC	15MG	A076901 001	Jun 28, 2005
	30MG	A076901 002	Jun 28, 2005
	45MG	A076901 003	Jun 28, 2005

MISOPROSTOLTABLET;ORAL
MISOPROSTOL

NOVEL LABS INC	0.1MG	A091667 001	Jul 25, 2012
	0.2MG	A091667 002	Jul 25, 2012

MITOMYCIN

INJECTABLE; INJECTION

MITOMYCIN

HIKMA	5MG/VIAL	A064117 001	Apr 19, 1995
HOSPIRA	20MG/VIAL	A064106 001	Nov 29, 1995

MITOZYTREX

+ SUPERGEN	5MG/VIAL **	N050763 001	Nov 14, 2002
------------	-------------	-------------	--------------

MUTAMYCIN

+ BRISTOL	5MG/VIAL **	N050450 001	
+ BRISTOL MYERS	20MG/VIAL **	N050450 002	
	5MG/VIAL	A062336 001	
	20MG/VIAL	A062336 002	
	40MG/VIAL **	A062336 003	Mar 10, 1988

MITOXANTRONE HYDROCHLORIDE

INJECTABLE; INJECTION

MITOXANTRONE HYDROCHLORIDE

FRESENIUS KABI ONCOL	EQ 20MG BASE/10ML (EQ 2MG BASE/ML)	A078606 001	May 14, 2008
	EQ 25MG BASE/12.5ML (EQ 2MG BASE/ML)	A078606 002	May 14, 2008
	EQ 30MG BASE/15ML (EQ 2MG BASE/ML)	A078606 003	May 14, 2008
RISING	EQ 20MG BASE/10ML (EQ 2MG BASE/ML)	A078980 001	Apr 13, 2009
	EQ 20MG BASE/10ML (EQ 2MG BASE/ML)	A201014 001	Dec 11, 2012
	EQ 30MG BASE/15ML (EQ 2MG BASE/ML)	A078980 002	Apr 13, 2009

NOVANTRONE

+ EMD SERONO	EQ 20MG BASE/10ML (EQ 2MG BASE/ML) **	N019297 001	Dec 23, 1987
+ EMD SERONO	EQ 25MG BASE/12.5ML (EQ 2MG BASE/ML) **	N019297 002	Dec 23, 1987
+ EMD SERONO	EQ 30MG BASE/15ML (EQ 2MG BASE/ML) **	N019297 003	Dec 23, 1987

MIVACURIUM CHLORIDE

INJECTABLE; INJECTION

MIVACRON IN DEXTROSE 5% IN PLASTIC CONTAINER

ABBVIE	EQ 0.5MG BASE/ML	N020098 002	Jan 22, 1992
	EQ 50MG BASE/100ML	N020098 003	Jan 22, 1992

MIVACURIUM CHLORIDE

RISING	EQ 2MG BASE/ML	A078562 001	Apr 30, 2009
--------	----------------	-------------	--------------

SOLUTION; INTRAVENOUS

MIVACRON

+ ABBVIE	EQ 2MG BASE/ML (EQ 2MG BASE/ML) **	N020098 001	Jan 22, 1992
+ ABBVIE	EQ 10MG BASE/5ML (EQ 2MG BASE/ML)	N020098 004	Jan 22, 1992
+ ABBVIE	EQ 20MG BASE/10ML (EQ 2MG BASE/ML)	N020098 005	Jan 22, 1992

MIVACURIUM CHLORIDE

WOODWARD	EQ 10MG BASE/5ML (EQ 2MG BASE/ML)	A209708 001	Oct 12, 2021
	EQ 20MG BASE/10ML (EQ 2MG BASE/ML)	A209708 002	Oct 12, 2021

MOBOCERTINIB SUCCINATE

CAPSULE; ORAL

EXKIVITY

+ TAKEDA PHARMS USA	EQ 40MG BASE	N215310 001	Sep 15, 2021
---------------------	--------------	-------------	--------------

MODAFINIL

TABLET; ORAL

MODAFINIL

HIKMA PHARMS	100MG	A090543 001	Sep 26, 2012
	200MG	A090543 002	Sep 26, 2012
NATCO PHARMA	100MG	A076594 001	Jul 16, 2012
	200MG	A076594 002	Jul 16, 2012

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

MOEXIPRIL HYDROCHLORIDE

TABLET; ORAL

UNIVASC

UCB INC

7.5MG **

N020312 001 Apr 19, 1995

15MG **

N020312 002 Apr 19, 1995

MOLINDONE HYDROCHLORIDE

CAPSULE; ORAL

MOBAN

+ CHARTWELL RX

5MG **

N017111 001

+

10MG **

N017111 002

+

25MG **

N017111 003

CONCENTRATE; ORAL

MOBAN

ENDO PHARMS

20MG/ML

N017938 001

TABLET; ORAL

MOBAN

+ CHARTWELL RX

5MG **

N017111 004

+

10MG **

N017111 005

+

25MG **

N017111 006

+

50MG **

N017111 007

+

100MG **

N017111 008

MOMETASONE FUROATE

CREAM; TOPICAL

ELOCON

ORGANON

0.1% **

N019625 001 May 06, 1987

+

0.1% **

N019625 002 Apr 19, 2013

MOMETASONE FUROATE

ANDA REPOSITORY

0.1%

A076591 001 Apr 18, 2007

FOUGERA PHARMS

0.1%

A076171 001 Apr 08, 2005

LOTION; TOPICAL

ELOCON

+ ORGANON

0.1% **

N019796 001 Mar 30, 1989

MOMETASONE FUROATE

COSETTE

0.1%

A077678 001 Nov 21, 2007

OINTMENT; TOPICAL

ELOCON

+ ORGANON

0.1%

N019543 001 Apr 30, 1987

MOMETASONE FUROATE

ENCUBE

0.1%

A076481 001 Nov 14, 2003

TARO

0.1%

A076624 001 Dec 03, 2004

SPRAY, METERED; NASAL

MOMETASONE FUROATE

AMNEAL

0.05MG/SPRAY

A217460 001 Jun 14, 2023

APOTEX

0.05MG/SPRAY

A217438 001 Apr 23, 2024

NASONEX

+ ORGANON LLC

0.05MG/SPRAY **

N020762 001 Oct 01, 1997

MONOBENZONE

CREAM; TOPICAL

BENOQUIN

VALEANT PHARM INTL

20%

N008173 003

MONOCTANOIN

LIQUID; PERFUSION, BILIARY

MOCTANIN

ETHITEK

100%

N019368 001 Oct 29, 1985

MONTELUKAST SODIUM

GRANULE; ORAL

MONTELUKAST SODIUM

NORVIUM BIOSCIENCE

EQ 4MG BASE/PACKET

A202776 001 Dec 18, 2012

TEVA PHARMS

EQ 4MG BASE/PACKET

A090955 001 Aug 03, 2012

TABLET; ORAL

MONTELUKAST SODIUM

AJANTA PHARMA LTD

EQ 10MG BASE

A203432 001 Jul 31, 2015

APOTEX CORP

EQ 10MG BASE

A201294 001 Aug 03, 2012

BRECKENRIDGE

EQ 10MG BASE

A205319 001 Oct 30, 2020

HIKMA

EQ 10MG BASE

A090655 001 Aug 03, 2012

L PERRIGO CO

EQ 10MG BASE

A206112 001 Apr 26, 2017

NORVIUM BIOSCIENCE

EQ 10MG BASE

A079103 001 Aug 03, 2012

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

MONTELUKAST SODIUM

TABLET;ORAL

MONTELUKAST SODIUM

SANDOZ INC	EQ 10MG BASE	A200889 001	Aug 03, 2012
STRIDES PHARMA	EQ 10MG BASE	A091576 001	Aug 03, 2012

TABLET, CHEWABLE;ORAL

MONTELUKAST SODIUM

AJANTA PHARMA LTD	EQ 4MG BASE	A203328 001	Jul 31, 2015
	EQ 5MG BASE	A203328 002	Jul 31, 2015
APOTEX INC	EQ 4MG BASE	A201508 001	Aug 03, 2012
	EQ 5MG BASE	A201508 002	Aug 03, 2012
HIKMA	EQ 4MG BASE	A091128 001	Aug 03, 2012
	EQ 5MG BASE	A091128 002	Aug 03, 2012
JUBILANT GENERICS	EQ 4MG BASE	A203795 001	Feb 27, 2015
	EQ 5MG BASE	A203795 002	Feb 27, 2015
NATCO PHARMA	EQ 4MG BASE	A079142 001	Aug 03, 2012
	EQ 5MG BASE	A079142 002	Aug 03, 2012
SANDOZ INC	EQ 4MG BASE	A091414 001	Aug 03, 2012
	EQ 5MG BASE	A091414 002	Aug 03, 2012
STRIDES PHARMA	EQ 4MG BASE	A091588 001	Aug 03, 2012
	EQ 5MG BASE	A091588 002	Aug 03, 2012
UNICHEM	EQ 4MG BASE	A208621 001	Jul 02, 2018
	EQ 5MG BASE	A208621 002	Jul 02, 2018

MORICIZINE HYDROCHLORIDE

TABLET;ORAL

ETHMOZINE

SHIRE	200MG	N019753 001	Jun 19, 1990
	250MG	N019753 002	Jun 19, 1990
	300MG	N019753 003	Jun 19, 1990

MORPHINE SULFATE

CAPSULE, EXTENDED RELEASE;ORAL

AVINZA

KING PHARMS LLC	30MG	N021260 001	Mar 20, 2002
	45MG	N021260 005	Dec 18, 2008
	60MG	N021260 002	Mar 20, 2002
	75MG	N021260 006	Dec 18, 2008
	90MG	N021260 003	Mar 20, 2002
	120MG	N021260 004	Mar 20, 2002

KADIAN

+	ALLERGAN	10MG	N020616 008	Apr 20, 2007
+		20MG	N020616 001	Jul 03, 1996
+		30MG	N020616 004	Mar 09, 2001
+		40MG	N020616 009	Jul 09, 2012
+		50MG	N020616 002	Jul 03, 1996
+		60MG	N020616 005	Mar 09, 2001
+		70MG	N020616 010	Jul 09, 2012
+		80MG	N020616 006	Oct 27, 2006
+		100MG	N020616 003	Jul 03, 1996
+		130MG	N020616 011	Jul 09, 2012
+		150MG	N020616 012	Jul 09, 2012
+		200MG	N020616 007	Feb 27, 2007

MORPHINE SULFATE

IMPAX LABS INC	20MG	A200411 001	Apr 12, 2016
	30MG	A200411 002	Apr 12, 2016
	40MG	A200411 007	Jul 25, 2018
	50MG	A200411 003	Apr 12, 2016
	60MG	A200411 004	Apr 12, 2016
	80MG	A200411 005	Apr 12, 2016
	100MG	A200411 006	Apr 12, 2016
NORTEC DEV ASSOC	20MG	A203158 001	Aug 04, 2021
	30MG	A203158 002	Aug 04, 2021
	50MG	A203158 003	Aug 04, 2021
	60MG	A203158 004	Aug 04, 2021
	80MG	A203158 005	Aug 04, 2021
	100MG	A203158 006	Aug 04, 2021
STRIDES PHARMA	20MG	A200812 001	Nov 10, 2011
	30MG	A200812 002	Nov 10, 2011
	50MG	A200812 003	Nov 10, 2011

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

MORPHINE SULFATECAPSULE, EXTENDED RELEASE;ORAL
MORPHINE SULFATE

	60MG	A200812 004	Nov 10, 2011
	80MG	A200812 005	Nov 10, 2011
	100MG	A200812 006	Nov 10, 2011
TEVA PHARMS USA	20MG	A202718 001	Dec 29, 2014
	30MG	A202718 002	Dec 29, 2014
	40MG	A202718 007	Jun 03, 2015
	50MG	A202718 003	Dec 29, 2014
	60MG	A202718 004	Dec 29, 2014
	70MG	A202718 008	Jun 03, 2015
	80MG	A202718 005	Dec 29, 2014
	100MG	A202718 006	Dec 29, 2014

INJECTABLE; INJECTION

ASTRAMORPH PF

FRESENIUS KABI USA	0.5MG/ML	A071050 001	Oct 07, 1986
	0.5MG/ML	A071051 001	Oct 07, 1986
	1MG/ML	A071052 001	Oct 07, 1986
	1MG/ML	A071053 001	Oct 07, 1986

MORPHINE SULFATE

HOSPIRA

	0.5MG/ML	A071849 001	May 11, 1988
	1MG/ML	A071850 001	May 11, 1988
+ HOSPIRA INC	15MG/ML **	N202515 005	Nov 14, 2011
ICU MEDICAL INC	0.5MG/ML	N019917 001	Oct 30, 1992
+	1MG/ML **	N019916 001	Oct 30, 1992
+	5MG/ML **	N019916 002	Oct 27, 2006
SPECGX LLC	1MG/ML	N020631 001	Jul 03, 1996
	2MG/ML	N020631 002	Jul 03, 1996
WATSON LABS	0.5MG/ML	A073373 001	Sep 30, 1991
	0.5MG/ML	A073375 001	Sep 30, 1991
	1MG/ML	A073374 001	Sep 30, 1991
	1MG/ML	A073376 001	Sep 30, 1991

INJECTABLE, LIPOSOMAL; EPIDURAL

DEPODUR

PACIRA PHARMS INC	10MG/ML (10MG/ML)	N021671 001	May 18, 2004
	15MG/1.5ML (10MG/ML)	N021671 002	May 18, 2004
	20MG/2ML (10MG/ML)	N021671 003	May 18, 2004

SOLUTION; INTRAMUSCULAR

MORPHINE SULFATE (AUTOINJECTOR)

+ MERIDIAN MEDCL TECHN	10MG/0.7ML (10MG/0.7ML)	N019999 001	Jul 12, 1990
------------------------	-------------------------	-------------	--------------

SOLUTION; INTRAMUSCULAR, INTRAVENOUS

MORPHINE SULFATE

+ FRESENIUS KABI USA	5MG/ML (5MG/ML)	N204223 003	Oct 30, 2013
+	8MG/ML (8MG/ML)	N204223 004	Oct 30, 2013

SOLUTION; ORAL

MORPHINE SULFATE

ANI PHARMS	10MG/5ML	A205509 001	Apr 17, 2018
	20MG/5ML	A205509 002	Apr 17, 2018
	100MG/5ML	A205509 003	Apr 17, 2018
CHARTWELL MOLECULAR	10MG/5ML	A202309 001	Nov 25, 2015
	20MG/5ML	A202310 001	Oct 30, 2015
	100MG/5ML	N201517 001	Jun 23, 2011
HIKMA	100MG/5ML	A208809 001	Jul 06, 2017
NOSTRUM LABS INC	10MG/5ML	A201011 001	Feb 05, 2014
	20MG/5ML	A201011 002	Feb 05, 2014
	100MG/5ML	A201011 003	Oct 06, 2016
PHARM ASSOC	10MG/5ML	A206573 002	Sep 12, 2023
	100MG/5ML	A206573 001	Nov 14, 2016
TRIS PHARMA INC	20MG/5ML	A203519 001	May 18, 2016
VISTAPHARM	10MG/5ML	A201947 001	Jan 05, 2012
	20MG/5ML	A201947 002	Jan 05, 2012

TABLET; ORAL

MORPHINE SULFATE

DR REDDYS LABS SA	15MG	A207270 001	Jan 12, 2022
	30MG	A207270 002	Jan 12, 2022
INGENUS PHARMS NJ	15MG	A215584 001	Feb 07, 2022
	30MG	A215584 002	Feb 07, 2022

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

MORPHINE SULFATE

TABLET, EXTENDED RELEASE;ORAL

ARYMO ER

+	ZYLA	15MG	N208603	001	Jan 09, 2017
+		30MG	N208603	002	Jan 09, 2017
+		60MG	N208603	003	Jan 09, 2017

MORPHABOND ER

+	OHEMO LIFE	15MG	N206544	001	Oct 02, 2015
+		30MG	N206544	002	Oct 02, 2015
+		60MG	N206544	003	Oct 02, 2015
+		100MG	N206544	004	Oct 02, 2015

MORPHINE SULFATE

	DR REDDYS LABS SA	15MG	A205386	001	Oct 28, 2016
		30MG	A205386	002	Oct 28, 2016
		60MG	A205386	003	Oct 28, 2016
		100MG	A205386	004	Oct 28, 2016
	EPIC PHARMA LLC	15MG	A091357	001	Jun 23, 2016
		30MG	A091357	002	Jun 23, 2016
		60MG	A091357	003	Jun 23, 2016
		100MG	A091357	004	Jun 23, 2016
		200MG	A091357	005	Jun 23, 2016
	NESHER PHARMS	15MG	A076733	001	May 19, 2004
		30MG	A076720	002	Dec 23, 2005
		60MG	A076720	001	May 19, 2004
		100MG	A077855	001	Sep 27, 2007
		200MG	A077855	002	Sep 27, 2007
	RISING	15MG	A200824	001	Oct 18, 2011
		30MG	A200824	002	Oct 18, 2011
		60MG	A200824	003	Oct 18, 2011
		100MG	A200824	004	Oct 18, 2011
		200MG	A200824	005	Oct 18, 2011
	SUN PHARM INDUSTRIES	15MG	A205634	001	Aug 25, 2016
		30MG	A205634	002	Aug 25, 2016
		60MG	A205634	003	Aug 25, 2016
		100MG	A205634	004	Aug 25, 2016
		200MG	A205634	005	Aug 25, 2016
	WATSON LABS	100MG	A075656	001	Jan 30, 2001
	ORAMORPH SR				
	XANODYNE PHARMS INC	15MG	N019977	004	Nov 23, 1994
		30MG	N019977	001	Aug 15, 1991
		60MG	N019977	002	Aug 15, 1991
		100MG	N019977	003	Aug 15, 1991

MORPHINE SULFATE; NALTREXONE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE;ORAL

EMBEDA

+	ALPHARMA PHARMS	20MG; 0.8MG **	N022321	001	Aug 13, 2009
+		30MG; 1.2MG **	N022321	002	Aug 13, 2009
+		50MG; 2MG **	N022321	003	Aug 13, 2009
+		60MG; 2.4MG **	N022321	004	Aug 13, 2009
+		80MG; 3.2MG **	N022321	005	Aug 13, 2009
+		100MG; 4MG **	N022321	006	Aug 13, 2009

MOXALACTAM DISODIUM

INJECTABLE; INJECTION

MOXAM

	LILLY	EQ 250MG BASE/VIAL	N050550	001	
		EQ 500MG BASE/VIAL	N050550	002	
		EQ 1GM BASE/VIAL	N050550	003	
		EQ 2GM BASE/VIAL	N050550	004	
		EQ 10GM BASE/VIAL	N050550	008	

MOXIFLOXACIN HYDROCHLORIDE

SOLUTION; INTRAVENOUS

AVELOX IN SODIUM CHLORIDE 0.8% IN PLASTIC CONTAINER

+	BAYER HLTHCARE	400MG/250ML (1.6MG/ML) **	N021277	001	Nov 30, 2001
---	----------------	---------------------------	---------	-----	--------------

SOLUTION/DROPS; OPHTHALMIC

MOXEZA

+	HARROW EYE	EQ 0.5% BASE	N022428	001	Nov 19, 2010
---	------------	--------------	---------	-----	--------------

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

MOXIFLOXACIN HYDROCHLORIDE

SOLUTION/DROPS;OPHTHALMIC

MOXIFLOXACIN HYDROCHLORIDE

EPIC PHARMA LLC EQ 0.5% BASE A202916 001 Nov 09, 2017

MYLAN EQ 0.5% BASE A206447 001 Mar 30, 2020

TABLET;ORAL

AVELOX

+ BAYER HLTHCARE EQ 400MG BASE ** N021085 001 Dec 10, 1999

MOXIFLOXACIN HYDROCHLORIDE

NATCO EQ 400MG BASE A204635 001 Aug 31, 2015

SUNSHINE EQ 400MG BASE A206295 001 Sep 28, 2018

TORRENT EQ 400MG BASE A200160 001 Apr 03, 2014

MUPIROICIN

OINTMENT;TOPICAL

BACTROBAN

+ GLAXOSMITHKLINE 2% ** N050591 001 Dec 31, 1987

MUPIROICIN CALCIUM

CREAM;TOPICAL

BACTROBAN

+ GLAXOSMITHKLINE EQ 2% BASE ** N050746 001 Dec 11, 1997

MUPIROICIN

PADAGIS ISRAEL EQ 2% BASE A212465 001 Aug 03, 2022

OINTMENT;NASAL

BACTROBAN

+ GLAXOSMITHKLINE EQ 2% BASE N050703 001 Sep 18, 1995

MYCOPHENOLATE MOFETIL

CAPSULE;ORAL

MYCOPHENOLATE MOFETIL

APOTEX CORP 250MG A090419 001 Apr 22, 2009

DR REDDYS LABS LTD 250MG A091315 001 Oct 27, 2011

JUBILANT CADISTA 250MG A090762 001 Dec 15, 2014

STRIDES PHARMA 250MG A090111 001 Dec 22, 2009

ZYDUS PHARMS USA INC 250MG A065433 001 May 04, 2009

TABLET;ORAL

MYCOPHENOLATE MOFETIL

AMNEAL 500MG A090606 001 Jul 16, 2010

APOTEX 500MG A090499 001 Apr 22, 2009

DR REDDYS LABS LTD 500MG A090464 001 Sep 13, 2010

JUBILANT CADISTA 500MG A090661 001 Dec 15, 2014

TEVA PHARMS 500MG A065457 001 May 04, 2009

ZYDUS PHARMS USA INC 500MG A065477 001 May 04, 2009

MYCOPHENOLATE MOFETIL HYDROCHLORIDE

INJECTABLE;INJECTION

MYCOPHENOLATE MOFETIL HYDROCHLORIDE

AMNEAL 500MG/VIAL A211374 001 Mar 05, 2021

MYCOPHENOLIC SODIUM

TABLET, DELAYED RELEASE;ORAL

MYCOPHENOLIC SODIUM

TEVA PHARMS USA EQ 180MG BASE A202720 001 Oct 30, 2014

EQ 360MG BASE A202720 002 Oct 30, 2014

NABILONE

CAPSULE;ORAL

CESAMET

+ BAUSCH 1MG N018677 001 Dec 26, 1985

NABUMETONE

TABLET;ORAL

NABUMETONE

AUROBINDO PHARMA USA 500MG A090516 001 Jul 12, 2010

750MG A090516 002 Jul 12, 2010

750MG A075179 001 Jun 06, 2000

750MG A075590 001 Feb 25, 2002

750MG A075590 002 Feb 25, 2002

750MG A075189 001 May 26, 2000

750MG A075189 002 Sep 24, 2001

750MG A090427 001 Dec 30, 2011

750MG A090427 002 Dec 30, 2011

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

NABUMETONE

TABLET; ORAL

NABUMETONE

OXFORD PHARMS

500MG

A079093 001 Feb 27, 2009

750MG

A079093 002 Feb 27, 2009

RELAFEN

+ SMITHKLINE BEECHAM

500MG **

N019583 001 Dec 24, 1991

+

750MG **

N019583 002 Dec 24, 1991

NADOLOL

TABLET; ORAL

CORCARD

+ USWM

20MG

N018063 005 Oct 28, 1986

+

40MG

N018063 001

+

80MG

N018063 002

120MG

N018063 003

160MG

N018063 004

NADOLOL

HERITAGE PHARMA

20MG

A074229 001 Aug 30, 1996

40MG

A074229 002 Aug 30, 1996

80MG

A074255 001 Jan 24, 1996

120MG

A074255 002 Jan 24, 1996

160MG

A074255 003 Jan 24, 1996

NOVAST LABS

20MG

A210786 001 Jun 01, 2018

40MG

A210786 002 Jun 01, 2018

80MG

A210786 003 Jun 01, 2018

TEVA PHARMS

80MG

A074368 001 Aug 31, 1994

120MG

A074368 002 Aug 31, 1994

160MG

A074368 003 Aug 31, 1994

NAFCILLIN SODIUM

CAPSULE; ORAL

UNIPEN

WYETH AYERST

EQ 250MG BASE

N050111 001

FOR SOLUTION; ORAL

UNIPEN

WYETH AYERST

EQ 250MG BASE/5ML

N050199 001

INJECTABLE; INJECTION

NAFCILLIN SODIUM

APOTHECON

EQ 500MG BASE/VIAL

A061984 001

EQ 1GM BASE/VIAL

A061984 002

EQ 2GM BASE/VIAL

A061984 003

EQ 4GM BASE/VIAL

A061984 005

FRESENIUS

EQ 1GM BASE/VIAL

A206682 001 Dec 10, 2019

EQ 2GM BASE/VIAL

A206682 002 Dec 10, 2019

SANDOZ

EQ 500MG BASE/VIAL

A062527 001 Aug 02, 1984

EQ 1GM BASE/VIAL

A062527 002 Aug 02, 1984

EQ 1GM BASE/VIAL

A062732 001 Dec 23, 1986

EQ 2GM BASE/VIAL

A062527 003 Aug 02, 1984

EQ 2GM BASE/VIAL

A062732 002 Dec 23, 1986

EQ 10GM BASE/VIAL

A062527 004 Aug 02, 1984

WATSON LABS INC

EQ 500MG BASE/VIAL

A062844 001 Oct 26, 1988

EQ 1GM BASE/VIAL

A062844 002 Oct 26, 1988

EQ 1.5GM BASE/VIAL

A062844 003 Oct 26, 1988

EQ 2GM BASE/VIAL

A062844 004 Oct 26, 1988

EQ 4GM BASE/VIAL

A062844 005 Oct 26, 1988

EQ 10GM BASE/VIAL

A063008 001 Sep 29, 1988

NALLPEN

GLAXOSMITHKLINE

EQ 500MG BASE/VIAL

A061999 001

EQ 1GM BASE/VIAL

A061999 002

EQ 1GM BASE/VIAL

A062755 001 Dec 19, 1986

EQ 2GM BASE/VIAL

A061999 003

EQ 2GM BASE/VIAL

A062755 002 Dec 19, 1986

EQ 10GM BASE/VIAL

A061999 004

UNIPEN

WYETH AYERST

EQ 500MG BASE/VIAL **

A062717 001 Dec 16, 1986

+

EQ 500MG BASE/VIAL **

N050320 001

EQ 1GM BASE/VIAL **

A062717 002 Dec 16, 1986

EQ 2GM BASE/VIAL **

A062717 004 Dec 16, 1986

+

EQ 2GM BASE/VIAL **

N050320 003

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

NAFCILLIN SODIUM

INJECTABLE; INJECTION

UNIPEN

+		EQ 4GM BASE/VIAL **	N050320	004
+		EQ 10GM BASE/VIAL **	N050320	005
+		EQ 20GM BASE/VIAL **	N050320	006

UNIPEN IN PLASTIC CONTAINER

+	WYETH AYERST	EQ 1GM BASE/VIAL **	N050320	002
---	--------------	---------------------	---------	-----

TABLET; ORAL

UNIPEN

	WYETH AYERST	EQ 500MG BASE	N050462	001
--	--------------	---------------	---------	-----

NAFTIFINE HYDROCHLORIDE

CREAM; TOPICAL

NAFTIN

+	LEGACY PHARMA	1% **	N019599	001	Feb 29, 1988
---	---------------	-------	---------	-----	--------------

NALBUPHINE HYDROCHLORIDE

INJECTABLE; INJECTION

NALBUPHINE

	ABRAXIS PHARM	10MG/ML	A070751	001	Jul 02, 1986
		20MG/ML	A070752	001	Sep 24, 1986

NALBUPHINE HYDROCHLORIDE

	ABBOTT	20MG/ML	A070917	001	Feb 03, 1989
	ABBVIE	1.5MG/ML	N020200	001	Mar 12, 1993
	DR REDDYS	10MG/ML	A074471	001	Mar 19, 1998
		20MG/ML	A074471	002	Mar 19, 1998
	IGI LABS INC	10MG/ML	A072070	001	Apr 10, 1989
		10MG/ML	A072071	001	Apr 10, 1989
		10MG/ML	A072072	001	Apr 10, 1989
		20MG/ML	A072073	001	Apr 10, 1989
		20MG/ML	A072074	001	Apr 10, 1989
		20MG/ML	A072075	001	Apr 10, 1989
	RISING	10MG/ML	A206506	001	Feb 06, 2019
		10MG/ML	A207595	001	Jan 11, 2019
		20MG/ML	A206506	002	Feb 06, 2019
		20MG/ML	A207595	002	Jan 11, 2019

NUBAIN

+	ENDO OPERATIONS	10MG/ML **	N018024	001	
+		20MG/ML **	N018024	002	May 27, 1982

NALIDIXIC ACID

SUSPENSION; ORAL

NEGGRAM

	SANOFI AVENTIS US	250MG/5ML	N017430	001
--	-------------------	-----------	---------	-----

TABLET; ORAL

NALIDIXIC ACID

	SUN PHARM INDUSTRIES	250MG	A070270	001	Jun 29, 1988
		500MG	A070271	001	Jun 29, 1988
		1GM	A070272	001	Jun 29, 1988
	WATSON LABS	250MG	A071936	001	Jun 29, 1988
		500MG	A072061	001	Jun 29, 1988
		1GM	A071919	001	Jun 29, 1988

NEGGRAM

	SANOFI AVENTIS US	250MG	N014214	002
		500MG	N014214	004
		1GM	N014214	005

NALMEFENE HYDROCHLORIDE

SOLUTION; INTRAMUSCULAR, INTRAVENOUS, SUBCUTANEOUS

REVEX

+	HIKMA	EQ 0.1MG BASE/ML (EQ 0.1MG BASE/ML) **	N020459	001	Apr 17, 1995
+		EQ 2MG BASE/2ML (EQ 1MG BASE/ML) **	N020459	002	Apr 17, 1995

NALOXONE HYDROCHLORIDE

INJECTABLE; INJECTION

NALOXONE

	HIKMA	0.4MG/ML	A070298	001	Sep 24, 1986
		0.4MG/ML	A070496	001	Sep 24, 1986
	WYETH AYERST	0.02MG/ML	A070188	001	Sep 24, 1986
		0.02MG/ML	A070189	001	Sep 24, 1986
		0.4MG/ML	A070190	001	Sep 24, 1986

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

NALOXONE HYDROCHLORIDE

INJECTABLE; INJECTION

NALOXONE

	0.4MG/ML	A070191 001	Sep 24, 1986
NALOXONE HYDROCHLORIDE			
ABRAXIS PHARM	0.02MG/ML	A070648 001	Nov 17, 1986
	0.02MG/ML	A070661 001	Nov 17, 1986
	0.4MG/ML	A070649 001	Nov 17, 1986
	1MG/ML	A071604 001	Dec 16, 1988
ASTRAZENECA	0.02MG/ML	A072081 001	Apr 11, 1989
CHARTWELL RX	0.4MG/ML	A207846 001	Dec 17, 2018
EUGIA PHARMA	0.4MG/ML	A212455 001	Oct 15, 2019
HIKMA	0.02MG/ML	A071272 001	May 24, 1988
	1MG/ML	A071273 001	May 24, 1988
	1MG/ML	A071274 001	May 24, 1988
	1MG/ML	A071287 001	May 24, 1988
HOSPIRA	0.02MG/ML	A070171 001	Sep 24, 1986
	0.02MG/ML	A070252 001	Jan 16, 1987
	0.02MG/ML	A070253 001	Jan 16, 1987
	0.4MG/ML	A070254 001	Jan 07, 1987
	0.4MG/ML	A070255 001	Jan 07, 1987
IGI LABS INC	0.02MG/ML	A072082 001	Apr 11, 1989
	0.02MG/ML	A072083 001	Apr 11, 1989
	0.02MG/ML	A072084 001	Apr 11, 1989
	0.02MG/ML	A072085 001	Apr 11, 1989
	0.4MG/ML	A072086 001	Apr 11, 1989
	0.4MG/ML	A072087 001	Apr 11, 1989
	0.4MG/ML	A072088 001	Apr 11, 1989
	0.4MG/ML	A072089 001	Apr 11, 1989
	0.4MG/ML	A072090 001	Apr 11, 1989
	1MG/ML	A072091 001	Apr 11, 1989
	1MG/ML	A072092 001	Apr 11, 1989
	1MG/ML	A072093 001	Apr 11, 1989
INTL MEDICATION	0.4MG/ML	A070417 001	Sep 24, 1986
	0.4MG/ML	A070639 001	Sep 24, 1986
	1MG/ML	A072115 001	Apr 27, 1988
MARSAM PHARMS LLC	0.4MG/ML	A071811 001	Jul 19, 1988
PAR STERILE PRODUCTS	0.4MG/ML	A211286 001	Jan 17, 2020
SMITH AND NEPHEW	0.02MG/ML	A071671 001	Nov 17, 1987
	0.4MG/ML	A071681 001	Nov 17, 1987
	0.4MG/ML	A071682 001	Nov 17, 1987
SOLOPAK	0.02MG/ML	A071672 001	Nov 17, 1987
	0.4MG/ML	A071683 001	Nov 17, 1987
WATSON LABS	0.4MG/ML	A071339 001	Nov 18, 1987
NARCAN			
+ ADAPT	0.02MG/ML **	N016636 002	
+	0.4MG/ML **	N016636 001	
+	1MG/ML **	N016636 003	Jun 14, 1982
BRISTOL MYERS SQUIBB	0.4MG/ML	A071083 001	Jul 28, 1988
	1MG/ML	A071084 001	Jul 28, 1988
	1MG/ML	A071311 001	Jul 28, 1988

SOLUTION; INTRAMUSCULAR, SUBCUTANEOUS

EVZIO

+ KALEO INC	0.4MG/0.4ML (0.4MG/0.4ML)	N205787 001	Apr 03, 2014
EVZIO (AUTOINJECTOR)			
+ KALEO INC	2MG/0.4ML (2MG/0.4ML)	N209862 001	Oct 19, 2016
NALOXONE HYDROCHLORIDE (AUTOINJECTOR)			
+ KALEO INC	10MG/0.4ML (10MG/0.4ML)	N215457 001	Feb 28, 2022

SPRAY, METERED; NASAL

NARCAN

+ EMERGENT	2MG/SPRAY **	N208411 002	Jan 24, 2017
------------	--------------	-------------	--------------

NALOXONE HYDROCHLORIDE; OXYCODONE HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL

TARGINIQ

+ PURDUE PHARMA LP	5MG; 10MG	N205777 001	Jul 23, 2014
+	10MG; 20MG	N205777 002	Jul 23, 2014
+	20MG; 40MG	N205777 003	Jul 23, 2014

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

NALOXONE HYDROCHLORIDE; PENTAZOCINE HYDROCHLORIDE

TABLET; ORAL

TALWIN NX

SANOFI AVENTIS US EQ 0.5MG BASE; EQ 50MG BASE ** N018733 001 Dec 16, 1982

NALTREXONE

FOR SUSPENSION, EXTENDED RELEASE; INTRAMUSCULAR

NALTREXONE

TEVA PHARMS USA INC 380MG/VIAL A213195 001 Jul 06, 2023

NALTREXONE HYDROCHLORIDE

TABLET; ORAL

NALTREXONE HYDROCHLORIDE

FOSUN PHARMA 50MG A075434 001 Mar 08, 2000

REVIA

+ TEVA WOMENS 50MG N018932 001 Nov 20, 1984

NALTREXONE HYDROCHLORIDE; OXYCODONE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

TROXYCA ER

PFIZER 1.2MG; 10MG N207621 001 Aug 19, 2016

2.4MG; 20MG N207621 002 Aug 19, 2016

3.6MG; 30MG N207621 003 Aug 19, 2016

4.8MG; 40MG N207621 004 Aug 19, 2016

7.2MG; 60MG N207621 005 Aug 19, 2016

9.6MG; 80MG N207621 006 Aug 19, 2016

NANDROLONE DECANOATE

INJECTABLE; INJECTION

DECA-DURABOLIN

WOODWARD SPECL 50MG/ML N013132 001 Jun 12, 1986

100MG/ML N013132 002 Jun 12, 1986

+ 200MG/ML ** N013132 003 Jun 12, 1986

NANDROLONE DECANOATE

ABRAXIS PHARM 100MG/ML A088290 001 Oct 03, 1983

200MG/ML A088317 001 Oct 14, 1983

AM REGENT 200MG/ML A091252 001 Aug 30, 2010

EPIC PHARMA LLC 100MG/ML A087519 001 Sep 28, 1983

WATSON LABS 50MG/ML A086385 001 Jan 13, 1984

50MG/ML A087598 001 Oct 06, 1983

50MG/ML A088554 001 Feb 10, 1986

100MG/ML A086598 001 Jan 13, 1984

100MG/ML A087599 001 Oct 06, 1983

+ 200MG/ML A088128 001 Dec 05, 1983

NANDROLONE PHENPROPIONATE

INJECTABLE; INJECTION

DURABOLIN

ORGANON USA INC 25MG/ML N011891 001

50MG/ML N011891 002

NANDROLONE PHENPROPIONATE

WATSON LABS 25MG/ML A086386 001 Jun 17, 1983

50MG/ML A087488 001 Jun 17, 1983

NAPHAZOLINE HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

ALBALON

ALLERGAN 0.1% ** A080248 001

NAFAZAIR

BAUSCH AND LOMB 0.1% A040073 001 May 25, 1994

PHARMAFAIR 0.1% A088101 001 Apr 15, 1983

NAPHAZOLINE HYDROCHLORIDE

RISING 0.1% A083590 001

NAPHCN FORTE

+ ALCON 0.1% A080229 001

OPCON

BAUSCH AND LOMB 0.1% A087506 001

VASOCON

NOVARTIS 0.1% A080235 002 Mar 24, 1983

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

NAPROXEN

TABLET; ORAL

NAPROSYN

+ ATNAHS PHARMA US

250MG

N017581 002

+

375MG

N017581 003

NAPROXEN

CHARTWELL MOLECULES

250MG

A074410 001 Apr 28, 1995

375MG

A074410 002 Apr 28, 1995

500MG

A074410 003 Apr 28, 1995

DAVA PHARMS INC

250MG

A074105 001 Dec 21, 1993

375MG

A074105 002 Dec 21, 1993

500MG

A074105 003 Dec 21, 1993

HAMILTON PHARMS

250MG

A074110 001 Oct 30, 1992

375MG

A074110 002 Oct 30, 1992

500MG

A074110 003 Oct 30, 1992

HIKMA INTL PHARMS

250MG

A076494 001 Jan 14, 2004

375MG

A076494 002 Jan 14, 2004

500MG

A076494 003 Jan 14, 2004

INVAGEN PHARMS

250MG

A091305 001 Aug 24, 2011

375MG

A091305 002 Aug 24, 2011

500MG

A091305 003 Aug 24, 2011

IVAX SUB TEVA PHARMS

250MG

A074111 001 Feb 28, 1995

375MG

A074111 002 Feb 28, 1995

500MG

A074111 003 Feb 28, 1995

L PERRIGO CO

250MG

A077339 001 Apr 27, 2005

375MG

A077339 002 Apr 27, 2005

500MG

A077339 003 Apr 27, 2005

NORVIUM BIOSCIENCE

250MG

A074121 001 Dec 21, 1993

375MG

A074121 002 Dec 21, 1993

500MG

A074121 003 Dec 21, 1993

PLIVA

250MG

A074182 001 Jun 27, 1996

375MG

A074182 002 Jun 27, 1996

500MG

A074182 003 Jun 27, 1996

PUREPAC PHARM

250MG

A074263 001 Dec 21, 1993

375MG

A074263 002 Dec 21, 1993

500MG

A074263 003 Dec 21, 1993

ROXANE

250MG

A074211 001 Feb 28, 1994

375MG

A074211 002 Feb 28, 1994

500MG

A074211 003 Feb 28, 1994

TEVA

250MG

A074129 001 Dec 21, 1993

250MG

A074216 001 Apr 11, 1996

375MG

A074129 002 Dec 21, 1993

375MG

A074216 002 Apr 11, 1996

500MG

A074129 003 Dec 21, 1993

500MG

A074216 003 Apr 11, 1996

TEVA PHARMS

250MG

A074207 001 Dec 21, 1993

375MG

A074207 002 Dec 21, 1993

500MG

A074207 003 Dec 21, 1993

WATSON LABS

250MG

A074457 001 May 31, 1995

375MG

A074457 002 May 31, 1995

500MG

A074457 003 May 31, 1995

WATSON LABS TEVA

250MG

A074163 001 Feb 10, 1995

375MG

A074163 002 Feb 10, 1995

500MG

A074163 003 Feb 10, 1995

TABLET, DELAYED RELEASE; ORAL

NAPROXEN

ACTAVIS ELIZABETH

375MG

A074936 001 Feb 24, 1998

500MG

A074936 002 Feb 24, 1998

AUROBINDO PHARMA USA

375MG

A075390 001 Apr 19, 2001

500MG

A075390 002 Apr 19, 2001

FOSUN PHARMA

375MG

A075061 001 Feb 18, 1998

500MG

A075061 002 Feb 18, 1998

PLIVA

375MG

A075337 001 May 26, 1999

500MG

A075337 002 May 26, 1999

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

NAPROXEN SODIUM

CAPSULE;ORAL

NAPROXEN SODIUM

STRIDES SOFTGELS

EQ 200MG BASE

A215472 001 Aug 17, 2022

TABLET;ORAL

ANAPROX

+ ATNAHS PHARMA US

EQ 250MG BASE **

N018164 001

NAPROXEN SODIUM

ABLE

EQ 250MG BASE

A076544 001 Aug 22, 2003

EQ 500MG BASE

A076544 002 Aug 22, 2003

AMNEAL PHARMS NY

EQ 250MG BASE

A078432 001 Apr 25, 2007

EQ 500MG BASE

A078432 002 Apr 25, 2007

CONTRACT PHARMACAL

220MG

A074789 001 Feb 27, 1997

HAMILTON PHARMS

EQ 250MG BASE

A074106 001 Aug 31, 1993

EQ 500MG BASE

A074106 002 Aug 31, 1993

HIKMA

EQ 250MG BASE

A074480 002 Feb 18, 1998

EQ 500MG BASE

A074480 001 May 14, 1996

IVAX SUB TEVA PHARMS

EQ 250MG BASE

A074230 001 Mar 14, 1995

EQ 500MG BASE

A074230 002 Mar 14, 1995

NORVIUM BIOSCIENCE

EQ 250MG BASE

A074367 001 Aug 31, 1994

EQ 500MG BASE

A074367 002 Aug 31, 1994

PLD ACQUISITIONS LLC

220MG

A074646 001 Jan 13, 1997

PLIVA

EQ 250MG BASE

A074242 001 Jun 20, 1996

EQ 500MG BASE

A074242 002 Jun 20, 1996

PUREPAC PHARM

EQ 250MG BASE

A074319 001 Mar 20, 1995

EQ 500MG BASE

A074319 002 Mar 20, 1995

ROXANE

EQ 250MG BASE

A074257 001 Dec 21, 1993

EQ 500MG BASE

A074257 002 Dec 21, 1993

SANDOZ

EQ 250MG BASE

A074162 001 Dec 21, 1993

EQ 250MG BASE

A074495 001 Dec 05, 1994

EQ 500MG BASE

A074162 002 Dec 21, 1993

EQ 500MG BASE

A074495 002 Dec 05, 1994

SUN PHARM INDS LTD

220MG

A091183 001 May 20, 2011

TEVA

EQ 250MG BASE

A074142 001 Dec 21, 1993

EQ 250MG BASE

A074198 001 Dec 21, 1993

EQ 500MG BASE

A074142 002 Dec 21, 1993

EQ 500MG BASE

A074198 002 Dec 21, 1993

TEVA PHARMS

EQ 250MG BASE

A074289 001 Jan 27, 1994

EQ 500MG BASE

A074289 002 Jan 27, 1994

WATSON LABS

EQ 250MG BASE

A074195 001 Dec 21, 1993

EQ 250MG BASE

A074455 001 May 31, 1995

EQ 500MG BASE

A074195 002 Dec 21, 1993

EQ 500MG BASE

A074455 002 May 31, 1995

NAPROXEN SODIUM; PSEUDOEPHEDRINE HYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL

ALEVE-D SINUS & COLD

+ BAYER

220MG;120MG **

N021076 001 Nov 29, 1999

NAPROXEN SODIUM; SUMATRIPTAN SUCCINATE

TABLET;ORAL

TREXIMET

+ CURRAX

60MG;EQ 10MG BASE

N021926 002 May 14, 2015

NARATRIPTAN HYDROCHLORIDE

TABLET;ORAL

AMERGE

+ GLAXOSMITHKLINE LLC

EQ 1MG BASE **

N020763 002 Feb 10, 1998

+

EQ 2.5MG BASE **

N020763 001 Feb 10, 1998

NARATRIPTAN

ANI PHARMS

EQ 1MG BASE

A078751 001 Jul 07, 2010

EQ 2.5MG BASE

A078751 002 Jul 07, 2010

APOTEX CORP

EQ 1MG BASE

A091373 001 Apr 22, 2011

EQ 2.5MG BASE

A091373 002 Apr 22, 2011

AUROBINDO PHARMA USA

EQ 1MG BASE

A202431 001 May 31, 2012

EQ 2.5MG BASE

A202431 002 May 31, 2012

CHARTWELL RX

EQ 1MG BASE

A090288 001 Jul 07, 2010

EQ 2.5MG BASE

A090288 002 Jul 07, 2010

SUN PHARM INDS LTD

EQ 2.5MG BASE

A091552 001 Feb 14, 2011

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

NATEGLINIDE

TABLET;ORAL

NATEGLINIDE

ALVOGEN	60MG	A205055 001	Dec 11, 2015
	120MG	A205055 002	Dec 11, 2015
TEVA PHARMS	60MG	A077467 001	Sep 09, 2009
	120MG	A077467 002	Sep 09, 2009
STARLIX			
+ NOVARTIS	60MG **	N021204 001	Dec 22, 2000
+	120MG **	N021204 002	Dec 22, 2000

NEBIVOLOL HYDROCHLORIDE

TABLET;ORAL

NEBIVOLOL HYDROCHLORIDE

ALKEM LABS LTD	EQ 2.5MG BASE	A203741 001	Jun 24, 2015
	EQ 5MG BASE	A203741 002	Jun 24, 2015
	EQ 10MG BASE	A203741 003	Jun 24, 2015
	EQ 20MG BASE	A203741 004	Jun 24, 2015
APOTEX	EQ 10MG BASE	A209741 001	Feb 27, 2024
	EQ 20MG BASE	A209741 002	Feb 27, 2024
CHARTWELL RX	EQ 2.5MG BASE	A213349 001	Mar 31, 2022
	EQ 5MG BASE	A213349 002	Mar 31, 2022
	EQ 10MG BASE	A213349 003	Mar 31, 2022
	EQ 20MG BASE	A213349 004	Mar 31, 2022
WATSON LABS INC	EQ 2.5MG BASE	A203683 001	Nov 27, 2015
	EQ 5MG BASE	A203683 002	Nov 27, 2015
	EQ 10MG BASE	A203683 003	Nov 27, 2015
	EQ 20MG BASE	A203683 004	Nov 27, 2015

NEBIVOLOL HYDROCHLORIDE; VALSARTAN

TABLET;ORAL

BYVALSON

+ ABBVIE	EQ 5MG BASE;80MG	N206302 001	Jun 03, 2016
----------	------------------	-------------	--------------

NEDOCROMIL SODIUM

AEROSOL, METERED; INHALATION

TILADE

KING PHARMS LLC	1.75MG/INH	N019660 001	Dec 30, 1992
-----------------	------------	-------------	--------------

SOLUTION; INHALATION

TILADE

SANOFI AVENTIS US	0.5%	N020750 001	Oct 01, 1997
-------------------	------	-------------	--------------

SOLUTION/DROPS; OPHTHALMIC

ALOCRIL

+ ALLERGAN	2% **	N021009 001	Dec 08, 1999
------------	-------	-------------	--------------

NEDOCROMIL SODIUM

EPIC PHARMA LLC	2%	A090638 001	Aug 22, 2012
-----------------	----	-------------	--------------

NEFAZODONE HYDROCHLORIDE

TABLET;ORAL

NEFAZODONE HYDROCHLORIDE

ANI PHARMS	50MG	A076072 001	Sep 16, 2003
	100MG	A076072 002	Sep 16, 2003
	150MG	A076072 003	Sep 16, 2003
	200MG	A076072 004	Sep 16, 2003
	250MG	A076072 005	Sep 16, 2003
AUROBINDO PHARMA USA	100MG	A076129 002	Sep 16, 2003
	150MG	A076129 003	Sep 16, 2003
	200MG	A076129 004	Sep 16, 2003
	250MG	A076129 005	Sep 16, 2003
CHARTWELL RX	50MG	A076302 001	Sep 16, 2003
	100MG	A076302 002	Sep 16, 2003
	150MG	A076302 003	Sep 16, 2003
	200MG	A076302 004	Sep 16, 2003
	250MG	A076302 005	Sep 16, 2003
DR REDDYS LABS INC	50MG	A076309 001	Sep 16, 2003
	100MG	A076309 002	Sep 16, 2003
	150MG	A076309 003	Sep 16, 2003
	200MG	A076309 004	Sep 16, 2003
	250MG	A076309 005	Sep 16, 2003
IVAX SUB TEVA PHARMS	50MG	A075763 001	Sep 16, 2003
	100MG	A075763 002	Sep 16, 2003

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

NEFAZODONE HYDROCHLORIDE

TABLET; ORAL

NEFAZODONE HYDROCHLORIDE

	150MG	A075763 003	Sep 16, 2003
	200MG	A075763 004	Sep 16, 2003
	250MG	A075763 005	Sep 16, 2003
ROXANE	50MG	A076196 001	Sep 16, 2003
	100MG	A076196 002	Sep 16, 2003
	150MG	A076196 003	Sep 16, 2003
	200MG	A076196 004	Sep 16, 2003
	250MG	A076196 005	Sep 16, 2003
SUN PHARM INDS LTD	50MG	A076409 001	Sep 16, 2003
	100MG	A076409 002	Sep 16, 2003
	150MG	A076409 003	Sep 16, 2003
	200MG	A076409 004	Sep 16, 2003
	250MG	A076409 005	Sep 16, 2003
WATSON LABS	100MG	A076073 002	Sep 16, 2003
	150MG	A076073 003	Sep 16, 2003
	200MG	A076073 004	Sep 16, 2003
	250MG	A076073 005	Sep 16, 2003
SERZONE			
+	BRISTOL MYERS SQUIBB 50MG **	N020152 001	Dec 22, 1994
+	100MG **	N020152 002	Dec 22, 1994
+	150MG **	N020152 003	Dec 22, 1994
+	200MG **	N020152 004	Dec 22, 1994
+	250MG **	N020152 005	Dec 22, 1994
+	300MG **	N020152 006	Dec 22, 1994

NELFINAVIR MESYLATE

POWDER; ORAL

VIRACEPT

AGOURON PHARMS EQ 50MG BASE/SCOOPFUL N020778 001 Mar 14, 1997

NEOMYCIN SULFATE

SOLUTION; ORAL

MYCIFRADIN

PHARMACIA AND UPJOHN EQ 87.5MG BASE/5ML N050285 001

NEO-FRADIN

X GEN PHARMS EQ 87.5MG BASE/5ML A065010 001 May 23, 2002

TABLET; ORAL

MYCIFRADIN

PHARMACIA AND UPJOHN EQ 350MG BASE A060520 001

NEOBIOTIC

PFIZER EQ 350MG BASE A060475 001

NEOMYCIN SULFATE

BRISTOL MYERS SQUIBB 500MG A060365 001

CHARTWELL MOLECULAR 500MG A204435 001 Jun 10, 2016

LANNETT 500MG A060607 001

LILLY 500MG A060385 001

NOSTRUM LABS INC 500MG A065468 001 Mar 29, 2010

ROXANE 500MG A062173 001

SANDOZ 500MG A061586 001

NEOMYCIN SULFATE; POLYMYXIN B SULFATE

CREAM; TOPICAL

NEOSPORIN

GLAXOSMITHKLINE EQ 3.5MG BASE/GM;10,000 UNITS/GM N050176 002 Jan 14, 1985

OINTMENT; OPHTHALMIC

STATROL

ALCON EQ 3.5MG BASE/GM;10,000 UNITS/GM N050344 002

SOLUTION/DROPS; OPHTHALMIC

STATROL

ALCON EQ 3.5MG BASE/ML;16,250 UNITS/ML A062339 001 Nov 30, 1984

EQ 3.5MG BASE/ML;16,250 UNITS/ML N050456 001

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

NEOMYCIN SULFATE; POLYMYXIN B SULFATE; PREDNISOLONE ACETATE

SUSPENSION/DROPS;OPHTHALMIC

POLY-PRED

ALLERGAN EQ 0.35% BASE;10,000 UNITS/ML;0.5% N050081 002

NEOMYCIN SULFATE; PREDNISOLONE ACETATE

OINTMENT;OPHTHALMIC

NEO-DELTA-CORTEF

PHARMACIA AND UPJOHN EQ 3.5MG BASE/GM;0.25% A061039 002

EQ 3.5MG BASE/GM;0.5% A061039 001

SUSPENSION/DROPS;OPHTHALMIC

NEO-DELTA-CORTEF

PHARMACIA AND UPJOHN EQ 3.5MG BASE/ML;0.25% A061037 001

NEOMYCIN SULFATE; PREDNISOLONE SODIUM PHOSPHATE

OINTMENT;OPHTHALMIC

NEO-HYDELTRASOL

MERCCK EQ 3.5MG BASE/GM;EQ 0.25% PHOSPHATE N050378 001

NEOMYCIN SULFATE; TRIAMCINOLONE ACETONIDE

CREAM;TOPICAL

MYTRES A

SAVAGE LABS EQ 3.5MG BASE/GM;0.1% A062598 001 Jul 21, 1986

NEOMYCIN SULFATE-TRIAMCINOLONE ACETONIDE

FOUGERA EQ 3.5MG BASE/GM;0.1% A062600 001 Jul 21, 1986

PHARMADERM EQ 3.5MG BASE/GM;0.1% A062595 001 Jul 21, 1986

OINTMENT;TOPICAL

MYTRES A

SAVAGE LABS EQ 3.5MG BASE/GM;0.1% A062609 001 May 23, 1986

NEOMYCIN SULFATE-TRIAMCINOLONE ACETONIDE

FOUGERA EQ 3.5MG BASE/GM;0.1% A062608 001 May 23, 1986

PHARMADERM EQ 3.5MG BASE/GM;0.1% A062607 001 May 23, 1986

NEOSTIGMINE METHYLSULFATE

SOLUTION;INTRAVENOUS

NEOSTIGMINE METHYLSULFATE

AM REGENT 5MG/10ML (0.5MG/ML) A209182 001 May 04, 2018

10MG/10ML (1MG/ML) A209182 002 May 04, 2018

EUGIA PHARMA 5MG/10ML (0.5MG/ML) A213244 001 Nov 02, 2023

10MG/10ML (1MG/ML) A213244 002 Nov 02, 2023

NESIRITIDE

FOR SOLUTION;INTRAVENOUS

NATRECOR

+ SCIOS LLC 1.5MG/VIAL ** N020920 001 Aug 10, 2001

NETILMICIN SULFATE

INJECTABLE;INJECTION

NETROMYCIN

SCHERING EQ 10MG BASE/ML N050544 001 Feb 28, 1983

EQ 25MG BASE/ML N050544 002 Feb 28, 1983

EQ 100MG BASE/ML N050544 003 Feb 28, 1983

NEVIRAPINE

SUSPENSION;ORAL

VIRAMUNE

+ BOEHRINGER INGELHEIM 50MG/5ML N020933 001 Sep 11, 1998

TABLET;ORAL

NEVIRAPINE

APOTEX INC 200MG A203021 001 May 22, 2012

NORVIUM BIOSCIENCE 200MG A078864 001 May 22, 2012

PRINSTON INC 200MG A078644 001 May 22, 2012

TECH ORGANIZED 200MG A203176 001 May 22, 2012

VIRAMUNE

+ BOEHRINGER INGELHEIM 200MG N020636 001 Jun 21, 1996

TABLET, EXTENDED RELEASE;ORAL

NEVIRAPINE

ALVOGEN 100MG A204621 002 Nov 09, 2015

400MG A204621 001 Jul 10, 2015

APOTEX 400MG A205258 001 Apr 03, 2014

AUROBINDO PHARMA 100MG A208616 001 Nov 23, 2016

400MG A207698 001 Feb 28, 2017

CIPLA 400MG A206448 001 Oct 15, 2015

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

NEVIRAPINE

TABLET, EXTENDED RELEASE;ORAL

NEVIRAPINE

NORVIUM BIOSCIENCE	100MG
TECH ORGANIZED	100MG
	400MG

A206271	001	Nov 09, 2015
A207467	001	Jul 31, 2017
A207467	002	Jul 31, 2017

VIRAMUNE XR

+ BOEHRINGER INGELHEIM	100MG
+	400MG

N201152	002	Nov 08, 2012
N201152	001	Mar 25, 2011

NIACIN

CAPSULE;ORAL

WAMPOCAP

MEDPOINTE PHARM HLC	500MG
---------------------	-------

N011073	003
---------	-----

TABLET;ORAL

NIACIN

EVERYLIFE	500MG
HALSEY	500MG
HIKMA	500MG
IMPAX LABS	500MG
IVAX SUB TEVA PHARMS	500MG
MK LABS	500MG
PUREPAC PHARM	500MG
SANDOZ	500MG
TABLICAPS	500MG
WATSON LABS	500MG
	500MG
	500MG
	500MG

A083203	001
A083453	001
A083718	001
A083115	001
A083180	001
A083525	001
A083271	001
A083306	001
A084237	001
A083136	001
A083305	001
A085172	001

WOCKHARDT

A081134	001	Apr 28, 1992
---------	-----	--------------

NICOLAR

+ SANOFI AVENTIS US	500MG
---------------------	-------

A083823	001
---------	-----

TABLET, EXTENDED RELEASE;ORAL

NIACIN

BEIJING	500MG
	1GM
JUBILANT GENERICS	500MG
	750MG
	1GM
RISING	500MG
	750MG
	1GM
YICHANG HUMANWELL	500MG
	750MG
	1GM

A214428	001	Nov 22, 2021
A214428	002	Nov 22, 2021
A209156	001	May 14, 2018
A209156	002	May 14, 2018
A209156	003	May 14, 2018
A203742	001	Feb 22, 2019
A203742	002	Feb 22, 2019
A203742	003	Feb 22, 2019
A212017	001	Jun 10, 2019
A212017	002	Jun 10, 2019
A212017	003	Jun 10, 2019

NIASPAN

ABBVIE	375MG
+	500MG **
+	750MG **
+	1GM **

N020381	001	Jul 28, 1997
N020381	002	Jul 28, 1997
N020381	003	Jul 28, 1997
N020381	004	Jul 28, 1997

NIASPAN TITRATION STARTER PACK

ABBVIE	375MG;500MG;750MG
--------	-------------------

N020381	005	Jul 28, 1997
---------	-----	--------------

NIACINAMIDE; PYRIDOXINE HYDROCHLORIDE; TYROSINE

SUSPENSION;ORAL

TPN

INTL MINERALS	15MG/5ML;3.75MG/5ML;600MG/5ML
---------------	-------------------------------

N008378	003
---------	-----

NICARDIPINE HYDROCHLORIDE

CAPSULE;ORAL

CARDENE

+ CHIESI	20MG **
+	30MG **

N019488	001	Dec 21, 1988
N019488	002	Dec 21, 1988

NICARDIPINE HYDROCHLORIDE

ANI PHARMS	20MG
	20MG
	30MG
	30MG
NORVIUM BIOSCIENCE	20MG
	30MG

A074439	001	Dec 10, 1996
A074540	001	Oct 28, 1996
A074439	002	Dec 10, 1996
A074540	002	Oct 28, 1996
A074642	001	Jul 18, 1996
A074642	002	Jul 18, 1996

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

NICARDIPINE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE;ORAL

CARDENE SR

+	CHIESI	30MG **	N020005 001	Feb 21, 1992
+		45MG **	N020005 002	Feb 21, 1992
+		60MG **	N020005 003	Feb 21, 1992

INJECTABLE;INJECTION

CARDENE

+	CHIESI	25MG/10ML (2.5MG/ML) **	N019734 001	Jan 30, 1992
---	--------	-------------------------	-------------	--------------

NICARDIPINE HYDROCHLORIDE

	HIKMA	25MG/10ML (2.5MG/ML)	A078714 001	Dec 28, 2009
	NAVINTA LLC	25MG/10ML (2.5MG/ML)	A090125 001	Nov 17, 2009
	SUN PHARM	25MG/10ML (2.5MG/ML)	N078405 001	Nov 17, 2009

INJECTABLE;INTRAVENOUS

CARDENE IN 5.0% DEXTROSE IN PLASTIC CONTAINER

+	CHIESI	40MG/200ML (0.2MG/ML)	N019734 005	Nov 07, 2008
---	--------	-----------------------	-------------	--------------

NICLOSAMIDE

TABLET, CHEWABLE;ORAL

NICLOCIDE

	BAYER PHARMS	500MG	N018669 001	May 14, 1982
--	--------------	-------	-------------	--------------

NICOTINE

FILM, EXTENDED RELEASE;TRANSDERMAL

NICOTROL

	MCNEIL CONS	15MG/16HR	N020536 001	Jul 03, 1996
--	-------------	-----------	-------------	--------------

PROSTEP

	AVEVA	11MG/24HR	N019983 003	Dec 23, 1998
		22MG/24HR	N019983 004	Dec 23, 1998

INHALANT;ORAL

NICOTROL

+	PFIZER	4MG/CARTRIDGE	N020714 001	May 02, 1997
---	--------	---------------	-------------	--------------

NICOTINE POLACRILEX

GUM, CHEWING;BUCCAL

NICOTINE POLACRILEX

	FERTIN PHARMA	EQ 4MG BASE	A214354 001	Dec 21, 2022
	IVAX SUB TEVA PHARMS	EQ 2MG BASE	A076880 001	Feb 18, 2009
		EQ 4MG BASE	A077850 001	Feb 18, 2009
	L PERRIGO CO	EQ 2MG BASE	A076776 001	Sep 16, 2004
		EQ 4MG BASE	A076778 001	Sep 16, 2004
	PERRIGO R AND D	EQ 2MG BASE	A078967 001	Apr 23, 2008
		EQ 4MG BASE	A078968 001	Apr 23, 2008

THRIVE

	GLAXOSMITHKLINE CONS	EQ 2MG BASE	A077658 001	Jun 19, 2007
		EQ 4MG BASE	A077656 001	Jun 19, 2007

NIFEDIPINE

CAPSULE;ORAL

ADALAT

	BAYER PHARMS	10MG	N019478 001	Nov 27, 1985
		20MG	N019478 002	Sep 17, 1986

NIFEDIPINE

	ACTAVIS ELIZABETH	20MG	A072556 001	Sep 20, 1990
	CHASE LABS NJ	10MG	A072409 001	Jul 04, 1990
		20MG	A073421 001	Jun 19, 1991
	TEVA	10MG	A072651 001	Feb 19, 1992

PROCARDIA

+	PFIZER	20MG **	N018482 002	Jul 24, 1986
---	--------	---------	-------------	--------------

TABLET, EXTENDED RELEASE;ORAL

ADALAT CC

+	NORWICH	30MG **	N020198 001	Apr 21, 1993
+		60MG **	N020198 002	Apr 21, 1993
+		90MG **	N020198 003	Apr 21, 1993

AFEDITAB CR

	WATSON LABS	60MG	A075659 001	Oct 26, 2001
	WATSON LABS TEVA	30MG	A075128 001	Mar 10, 2000

NIFEDIPINE

	AUROBINDO PHARMA USA	30MG	A090649 001	Jun 21, 2010
		60MG	A090649 002	Jun 21, 2010
		90MG	A090649 003	Jun 21, 2010

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

NIFEDIPINE

TABLET, EXTENDED RELEASE;ORAL

NIFEDIPINE

ENDO OPERATIONS	30MG	A077899 001	Dec 13, 2006
	60MG	A077899 002	Dec 13, 2006
	90MG	A077899 003	May 25, 2012
MARTEC USA LLC	90MG	A075414 003	Mar 23, 2004
NORVIUM BIOSCIENCE	30MG	A075108 001	Dec 17, 1999
RISING	30MG	A090602 001	Sep 13, 2010
	30MG	A201071 001	Dec 03, 2010
	60MG	A090602 002	Sep 13, 2010
	60MG	A201071 002	Dec 03, 2010
	90MG	A090602 003	Sep 13, 2010
	90MG	A201071 003	Dec 03, 2010
SWISS PHARM	30MG	A216019 001	Nov 18, 2022
	60MG	A216019 002	Nov 18, 2022

NILOTINIB HYDROCHLORIDE

CAPSULE;ORAL

NILOTINIB HYDROCHLORIDE

APOTEX	EQ 50MG BASE	A203640 001	Jan 05, 2024
	EQ 150MG BASE	A203640 002	Jan 05, 2024
	EQ 200MG BASE	A203640 003	Jan 05, 2024

NILUTAMIDE

TABLET;ORAL

NILANDRON

ADVANZ PHARMA	50MG	N020169 001	Sep 19, 1996
---------------	------	-------------	--------------

NIMODIPINE

CAPSULE;ORAL

NIMODIPINE

SOFGEN PHARMS	30MG	A201832 001	Jul 24, 2015
SUN PHARM INDS INC	30MG	A077067 001	Apr 17, 2007

NIMOTOP

+ BAYER PHARMS	30MG **	N018869 001	Dec 28, 1988
----------------	---------	-------------	--------------

SOLUTION;ORAL

NYMALIZE

+ AZURITY	3MG/ML **	N203340 001	May 10, 2013
-----------	-----------	-------------	--------------

NIRAPARIB TOSYLATE

CAPSULE;ORAL

ZEJULA

+ GLAXOSMITHKLINE	EQ 100MG BASE	N208447 001	Mar 27, 2017
-------------------	---------------	-------------	--------------

NISOLDIPINE

TABLET, EXTENDED RELEASE;ORAL

NISOLDIPINE

AMTA	8.5MG	A216606 001	Apr 10, 2023
------	-------	-------------	--------------

SULAR

+ COVIS	10MG **	N020356 001	Feb 02, 1995
---------	---------	-------------	--------------

	20MG **	N020356 002	Feb 02, 1995
--	---------	-------------	--------------

	25.5MG **	N020356 006	Jan 02, 2008
--	-----------	-------------	--------------

	30MG **	N020356 003	Feb 02, 1995
--	---------	-------------	--------------

	40MG **	N020356 004	Feb 02, 1995
--	---------	-------------	--------------

NITAZOXANIDE

FOR SUSPENSION;ORAL

ALINIA

+ ROMARK	100MG/5ML	N021498 001	Nov 22, 2002
----------	-----------	-------------	--------------

TABLET;ORAL

ALINIA

+ ROMARK	500MG	N021497 001	Jul 21, 2004
----------	-------	-------------	--------------

NITISINONE

CAPSULE;ORAL

NITISINONE

TORRENT	2MG	A215908 001	Jan 09, 2023
	5MG	A215908 002	Jan 09, 2023
	10MG	A215908 003	Jan 09, 2023
	20MG	A215908 004	Jan 09, 2023

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

NITRIC OXIDE

GAS; INHALATION

INOMAX

+ MALLINCKRODT HOSP 100PPM ** N020845 002 Dec 23, 1999

NITROFURANTOIN

CAPSULE; ORAL

NITROFURANTOIN

WATSON LABS 50MG A084326 001

100MG A084326 002

TABLET; ORAL

FURADANTIN

PROCTER AND GAMBLE 50MG N008693 001

100MG N008693 002

FURALAN

CHARTWELL MOLECULAR 50MG A080017 001

100MG A080017 002

NITROFURANTOIN

ELKINS SINN 50MG A080003 001

100MG A080003 002

IVAX SUB TEVA PHARMS 50MG A080078 002

100MG A080078 001

SANDOZ 50MG A080043 001

100MG A080043 002

WATSON LABS 50MG A080447 001

50MG A085797 001

100MG A080447 002

100MG A085796 001

WHITEWORTH TOWN PLSN 100MG A084085 002

NITROFURANTOIN SODIUM

INJECTABLE; INJECTION

IVADANTIN

PROCTER AND GAMBLE EQ 180MG BASE/VIAL N012402 001

NITROFURANTOIN, MACROCRYSTALLINE

CAPSULE; ORAL

NITROFURANTOIN

ACTAVIS LABS FL INC 25MG A091095 001 Jun 18, 2015

50MG A091095 002 Jun 18, 2015

100MG A091095 003 Jun 18, 2015

ATHEM 25MG A074336 001 Jan 25, 1995

50MG A074336 002 Jan 25, 1995

100MG A074336 003 Jan 25, 1995

WATSON LABS 25MG A073696 001 Dec 31, 1992

50MG A073696 002 Dec 31, 1992

100MG A073696 003 Dec 31, 1992

NITROFURANTOIN MACROCRYSTALLINE

WATSON LABS 50MG A070248 001 Jun 24, 1988

100MG A070249 001 Jun 24, 1988

NITROFURANTOIN; NITROFURANTOIN, MACROCRYSTALLINE

CAPSULE; ORAL

NITROFURANTOIN (MONOHYDRATE/MACROCRYSTALS)

ALVOGEN 75MG; 25MG A215002 001 Jul 20, 2022

CHARTWELL RX 75MG; 25MG A076648 001 Mar 22, 2004

RANBAXY LABS LTD 75MG; 25MG A076951 001 Mar 30, 2005

NITROFURAZONE

CREAM; TOPICAL

FURACIN

SHIRE 0.2% A083789 001

DRESSING; TOPICAL

ACTIN-N

SHERWOOD MEDCL 0.2% N017343 001

OINTMENT; TOPICAL

FURACIN

SHIRE 0.2% N005795 001

NITROFURAZONE

AMBIX 0.2% A086077 001

LANNETT 0.2% A084393 001

PERRIGO NEW YORK 0.2% A084968 001

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

NITROFURAZONE

OINTMENT; TOPICAL

NITROFURAZONE

TARO	0.2%	A086156	001
WENDT	0.2%	A086766	001

POWDER; TOPICAL

FURACIN

SHIRE	0.2%	A083791	001
-------	------	---------	-----

SOLUTION; TOPICAL

NITROFURAZONE

PERRIGO NEW YORK	0.2%	A085130	001
WENDT	0.2%	A087081	001

NITROGLYCERIN

AEROSOL; SUBLINGUAL

NITROLINGUAL

POHL BOSKAMP	0.4MG/SPRAY	N018705	001	Oct 31, 1985
--------------	-------------	---------	-----	--------------

FILM, EXTENDED RELEASE; TRANSDERMAL

MINITRAN

BAUSCH	0.4MG/HR	A089773	001	Aug 30, 1996
VALEANT PHARMS	0.1MG/HR	A089771	001	Aug 30, 1996
	0.6MG/HR	A089774	001	Aug 30, 1996
VALEANT PHARMS NORTH	0.2MG/HR	A089772	001	Aug 30, 1996

NITROGLYCERIN

LANNETT CO INC	0.2MG/HR	A075115	001	Aug 10, 2004
	0.4MG/HR	A075115	002	Aug 10, 2004
MYLAN TECHNOLOGIES	0.1MG/HR	A074992	004	Nov 12, 1999
	0.2MG/HR	A074992	003	Nov 12, 1999
	0.4MG/HR	A074992	002	Nov 12, 1999
	0.6MG/HR	A074992	001	Nov 12, 1999

TRANSDERM-NITRO

+ NOVARTIS	0.1MG/HR **	N020144	001	Feb 27, 1996
+	0.2MG/HR **	N020144	002	Feb 27, 1996
+	0.4MG/HR **	N020144	003	Feb 27, 1996
+	0.6MG/HR **	N020144	004	Feb 27, 1996
+	0.8MG/HR **	N020144	005	Feb 27, 1996

INJECTABLE; INJECTION

NITRO IV

POHL BOSKAMP	5MG/ML	N018672	002	Aug 30, 1983
--------------	--------	---------	-----	--------------

NITRO-BID

SANOFI AVENTIS US	5MG/ML	N018621	001	Jan 05, 1982
	10MG/ML	A071159	001	Feb 28, 1990

NITROGLYCERIN

ABRAXIS PHARM	5MG/ML	A070077	001	Dec 13, 1985
	5MG/ML	A071203	001	May 08, 1987
+ HOSPIRA	5MG/ML **	N018531	001	
INTL MEDICATION	5MG/ML	A070026	001	Sep 10, 1985
LUITPOLD	5MG/ML	A071492	001	May 24, 1988
SMITH AND NEPHEW	5MG/ML	A070633	001	Jun 19, 1986
	5MG/ML	A070634	001	Jun 19, 1986

NITROGLYCERIN IN DEXTROSE 5%

HOSPIRA	0.1MG/ML	A074083	001	Oct 26, 1994
	10MG/100ML	A071846	001	Aug 31, 1990
	20MG/100ML	A071847	001	Aug 31, 1990
	40MG/100ML	A071848	001	Aug 31, 1990

NITROL

RORER	0.8MG/ML	N018774	001	Jan 19, 1983
-------	----------	---------	-----	--------------

NITRONAL

POHL BOSKAMP	1MG/ML	N018672	001	Aug 30, 1983
--------------	--------	---------	-----	--------------

NITROSTAT

PARKE DAVIS	0.8MG/ML	N018588	001	
	5MG/ML	A070863	001	Jan 08, 1987
	5MG/ML	N018588	002	Dec 23, 1983
	10MG/ML	A070871	001	Jan 08, 1987
	10MG/ML	A070872	001	Jan 08, 1987

TRIDIL

HOSPIRA	0.5MG/ML	N018537	002	Jun 16, 1983
	5MG/ML	N018537	001	

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

NITROGLYCERIN

POWDER; SUBLINGUAL

GONITRO

+ POHL BOSKAMP

0.4MG/PACKET

N208424 001 Jun 08, 2016

TABLET; SUBLINGUAL

NITROGLYCERIN

ACTAVIS LABS FL INC

0.3MG

A203693 001 Oct 16, 2017

0.4MG

A203693 002 Oct 16, 2017

0.6MG

A203693 003 Oct 16, 2017

SIGMAPHARM LABS LLC

0.3MG

A207745 001 May 07, 2018

0.4MG

A207745 002 May 07, 2018

0.6MG

A207745 003 May 07, 2018

ZYDUS PHARMS

0.3MG

A210153 001 Mar 08, 2022

0.4MG

A210153 002 Mar 08, 2022

0.6MG

A210153 003 Mar 08, 2022

NIZATIDINE

CAPSULE; ORAL

AXID

SMITHKLINE BEECHAM

150MG

N019508 001 Apr 12, 1988

300MG

N019508 002 Apr 12, 1988

NIZATIDINE

ANI PHARMS

150MG

A075461 001 Jul 08, 2002

150MG

A075668 001 Sep 12, 2002

300MG

A075461 002 Jul 08, 2002

300MG

A075668 002 Sep 12, 2002

APOTEX INC

150MG

A076383 001 Jan 23, 2003

300MG

A076383 002 Jan 23, 2003

MYLAN PHARMS INC

150MG

A075934 001 Jul 09, 2002

300MG

A075934 002 Jul 09, 2002

NORVIUM BIOSCIENCE

150MG

A075806 001 Jul 05, 2002

300MG

A075806 002 Jul 05, 2002

SOLUTION; ORAL

AXID

+ BRAINTREE

15MG/ML **

N021494 001 May 25, 2004

NIZATIDINE

AMNEAL PHARMS

15MG/ML

A090576 001 Nov 18, 2009

NONOXYNOL-9

AEROSOL; VAGINAL

DELFIN

PERSONAL PRODS

12.5%

N014349 002

SPONGE; VAGINAL

TODAY

+ MAYER LABS INC

1GM

N018683 001 Apr 01, 1983

NOREPINEPHRINE BITARTRATE

INJECTABLE; INJECTION

NOREPINEPHRINE BITARTRATE

METRICS PHARM

EQ 1MG BASE/ML

A040522 001 Sep 30, 2004

SUN PHARM

EQ 1MG BASE/ML

A211980 001 Jan 29, 2021

ZYDUS PHARMS

EQ 1MG BASE/ML

A216341 001 Jul 19, 2022

NOREPINEPHRINE BITARTRATE; PROCAINE HYDROCHLORIDE; PROPOXYCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

RAVOCAINE AND NOVOCAIN W/ LEVOPHED

EASTMAN KODAK

EQ 0.033MG BASE/ML; 2%; 0.4%

N008592 003

NORETHINDRONE

TABLET; ORAL

NORLUTIN

PARKE DAVIS

5MG

N010895 002

TABLET; ORAL-28

MICRONOR

+ JANSSEN PHARMS

0.35MG **

N016954 001

NORETHINDRONE

AMNEAL PHARMS

0.35MG

A202260 001 Aug 01, 2013

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

NORETHINDRONE ACETATE

TABLET; ORAL

AYGESTIN

+ DURAMED RES 5MG ** N018405 001 Apr 21, 1982

NORETHINDRONE ACETATE

AUROBINDO PHARMA LTD 5MG A204236 001 Jan 08, 2016

NORLUTATE

PARKE DAVIS 5MG N012184 002

NORFLOXACIN

SOLUTION/DROPS; OPHTHALMIC

CHIBROXIN

MERCK 0.3% N019757 001 Jun 17, 1991

TABLET; ORAL

NOROXIN

+ MERCK 400MG ** N019384 002 Oct 31, 1986

NORTRIPTYLINE HYDROCHLORIDE

CAPSULE; ORAL

AVENTYL HYDROCHLORIDE

LILLY EQ 10MG BASE N014684 001

EQ 25MG BASE N014684 002

NORTRIPTYLINE HYDROCHLORIDE

ANI PHARMS EQ 10MG BASE A074054 001 Dec 31, 1992

EQ 25MG BASE A074054 002 Dec 31, 1992

EQ 50MG BASE A074054 003 Dec 31, 1992

EQ 75MG BASE A074054 004 Dec 31, 1992

AUROBINDO PHARMA LTD EQ 10MG BASE A074835 001 Jun 30, 1997

EQ 25MG BASE A074835 002 Jun 30, 1997

EQ 50MG BASE A074835 003 Jun 30, 1997

EQ 75MG BASE A074835 004 Jun 30, 1997

RISING EQ 10MG BASE A074234 001 Jul 26, 1993

EQ 25MG BASE A074234 002 Jul 26, 1993

EQ 50MG BASE A074234 003 Jul 26, 1993

EQ 75MG BASE A074234 004 Jul 26, 1993

TEVA EQ 10MG BASE A073667 001 Apr 11, 1996

EQ 25MG BASE A073667 002 Apr 11, 1996

EQ 50MG BASE A073667 003 Apr 11, 1996

EQ 75MG BASE A073667 004 Apr 11, 1996

ZYDUS LIFESCIENCES EQ 10MG BASE A213441 001 Feb 24, 2021

EQ 25MG BASE A213441 002 Feb 24, 2021

EQ 50MG BASE A213441 003 Feb 24, 2021

EQ 75MG BASE A213441 004 Feb 24, 2021

PAMELOR

+ SPECGX LLC EQ 10MG BASE N018013 001

+ EQ 25MG BASE N018013 002

+ EQ 50MG BASE N018013 004

+ EQ 75MG BASE N018013 003

SOLUTION; ORAL

AVENTYL

+ RANBAXY EQ 10MG BASE/5ML ** N014685 001

NORTRIPTYLINE HYDROCHLORIDE

TARO EQ 10MG BASE/5ML A077965 001 Jun 20, 2006

PAMELOR

SPECGX LLC EQ 10MG BASE/5ML N018012 001

NYSTATIN

CREAM; TOPICAL

CANDEX

BAYER PHARMS 100,000 UNITS/GM A061810 001

MYCOSTATIN

DELCOR ASSET CORP 100,000 UNITS/GM ** A060575 001

MYKINAC

ALPHARMA US PHARMS 100,000 UNITS/GM A062387 001 Jul 29, 1982

NILSTAT

LEDERLE 100,000 UNITS/GM A061445 001

NYSTATIN

STRIDES PHARMA 100,000 UNITS/GM A065315 001 May 31, 2006

TARO 100,000 UNITS/GM A062457 001 Jul 28, 1983

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

NYSTATIN

LOTION; TOPICAL					
CANDEX					
BAYER PHARMS	100,000 UNITS/ML			N050233 001	
OINTMENT; TOPICAL					
MYCOSTATIN					
DELCOR ASSET CORP	100,000 UNITS/GM **			A060571 001	
MYKINAC					
ALPHARMA US PHARMS	100,000 UNITS/GM			A062731 001	Sep 22, 1986
NILSTAT					
LEDERLE	100,000 UNITS/GM			A061444 001	
PASTILLE; ORAL					
MYCOSTATIN					
DELCOR ASSET CORP	200,000 UNITS			N050619 001	Apr 09, 1987
POWDER; ORAL					
BARSTATIN 100					
BARLAN	100%			A062489 001	Apr 27, 1988
NILSTAT					
+ STRIDES PHARMA	100% **			N050576 001	Dec 22, 1983
NYSTATIN					
PADDOCK LLC	100%			A062613 001	Nov 26, 1985
POWDER; TOPICAL					
MYCOSTATIN					
DELCOR ASSET CORP	100,000 UNITS/GM **			A060578 001	
NYSTATIN					
NESHER PHARMS	100,000 UNITS/GM			A065321 001	Aug 18, 2006
SUPPOSITORY; VAGINAL					
NYSERT					
WARNER CHILCOTT	100,000 UNITS			N050478 001	
SUSPENSION; ORAL					
MYCOSTATIN					
DELCOR ASSET CORP	100,000 UNITS/ML			A061533 001	
NILSTAT					
+ CHARTWELL MOLECULES	100,000 UNITS/ML **			N050299 001	
NYSTATIN					
ALPHARMA US PHARMS	100,000 UNITS/ML			A062571 001	Oct 29, 1985
COSETTE	100,000 UNITS/ML			A062776 001	Dec 17, 1987
G AND W LABS INC	100,000 UNITS/ML			A062349 001	Jul 14, 1982
MLV	100,000 UNITS/ML			A062832 001	Dec 27, 1991
MORTON GROVE	100,000 UNITS/ML			A062835 001	Nov 19, 1987
PAI HOLDINGS	100,000 UNITS/ML			A064042 001	Feb 28, 1994
PHARMADERM	100,000 UNITS/ML			A062518 001	Jul 06, 1984
PHARMAFAIR	100,000 UNITS/ML			A062541 001	Jan 16, 1985
TEVA	100,000 UNITS/ML			A062670 001	Jun 18, 1987
VISTAPHARM LLC	100,000 UNITS/ML			A064142 001	Jun 25, 1998
	100,000 UNITS/ML			A064142 002	Mar 07, 2011
NYSTEX					
SAVAGE LABS	100,000 UNITS/ML			A062519 001	Jul 06, 1984
TABLET; ORAL					
MYCOSTATIN					
DELCOR ASSET CORP	500,000 UNITS			A060574 001	
NILSTAT					
LEDERLE	500,000 UNITS			A061151 001	
NYSTATIN					
CHARTWELL RX	500,000 UNITS			A062524 001	Nov 26, 1985
QUANTUM PHARMICS	500,000 UNITS			A062525 001	Oct 29, 1984
SANDOZ	500,000 UNITS			A062065 001	
SUN PHARM INDUSTRIES	500,000 UNITS			A062838 001	Dec 22, 1988
WATSON LABS	500,000 UNITS			A062402 001	Dec 16, 1982
TABLET; VAGINAL					
KOROSTATIN					
HOLLAND RANTOS	100,000 UNITS			A061718 001	
MYCOSTATIN					
DELCOR ASSET CORP	100,000 UNITS			A060577 001	
NILSTAT					
LEDERLE	100,000 UNITS			A061325 001	
NYSTATIN					
FOUGERA	100,000 UNITS			A062459 001	Nov 09, 1983
ODYSSEY PHARMS	100,000 UNITS			A062615 001	Oct 17, 1985

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

NYSTATIN

TABLET; VAGINAL

NYSTATIN

PHARMADERM	100,000 UNITS	A062460 001	Nov 09, 1983
QUANTUM PHARMICS	100,000 UNITS	A062509 001	Apr 03, 1984
SANDOZ	100,000 UNITS	A061965 001	
TEVA	100,000 UNITS	A062502 001	Dec 23, 1983
WATSON LABS	100,000 UNITS	A062176 001	

NYSTATIN; TRIAMCINOLONE ACETONIDE

CREAM; TOPICAL

MYCO-TRIA CET II

TEVA	100,000 UNITS/GM;0.1%	A061954 002	Sep 20, 1985
------	-----------------------	-------------	--------------

MYCOLOG-II

DEL COR ASSET CORP	100,000 UNITS/GM;0.1% **	A060576 002	May 01, 1985
MYLAN	100,000 UNITS/GM;0.1% **	A062606 001	May 15, 1985

MYKACET

COSETTE	100,000 UNITS/GM;0.1%	A062367 001	May 28, 1985
---------	-----------------------	-------------	--------------

MYTRES F

SAVAGE LABS	100,000 UNITS/GM;0.1%	A062597 001	Oct 08, 1985
-------------	-----------------------	-------------	--------------

NYSTATIN AND TRIAMCINOLONE ACETONIDE

ALPHARMA US PHARMS	100,000 UNITS/GM;0.1%	A063010 001	Dec 20, 1988
PERRIGO NEW YORK	100,000 UNITS/GM;0.1%	A062186 002	Jun 06, 1985
PHARMAFAIR	100,000 UNITS/GM;0.1%	A062657 001	Jul 30, 1986
TARO	100,000 UNITS/GM;0.1%	A062347 001	Mar 30, 1987
TORRENT	100,000 UNITS/GM;0.1%	A213142 001	Jul 14, 2020

NYSTATIN TRIAMCINOLONE ACETONIDE

PHARMADERM	100,000 UNITS/GM;0.1%	A062596 001	Oct 08, 1985
------------	-----------------------	-------------	--------------

OINTMENT; TOPICAL

MYCO-TRIA CET II

TEVA	100,000 UNITS/GM;0.1%	A062045 002	Nov 26, 1985
------	-----------------------	-------------	--------------

MYCOLOG-II

MYLAN	100,000 UNITS/GM;0.1% **	A060572 001	Jun 28, 1985
-------	--------------------------	-------------	--------------

MYKACET

COSETTE	100,000 UNITS/GM;0.1%	A062733 001	Mar 06, 1987
---------	-----------------------	-------------	--------------

MYTRES F

SAVAGE LABS	100,000 UNITS/GM;0.1%	A062601 001	Oct 09, 1985
-------------	-----------------------	-------------	--------------

NYSTATIN AND TRIAMCINOLONE ACETONIDE

CHARTWELL RX	100,000 UNITS/GM;0.1%	A207316 001	Nov 18, 2019
CROWN LABS INC	100,000 UNITS/GM;0.1%	A207731 001	Dec 26, 2017
PAI HOLDINGS PHARM	100,000 UNITS/GM;0.1%	A208287 001	Dec 30, 2016
PERRIGO NEW YORK	100,000 UNITS/GM;0.1%	A062280 002	Oct 10, 1985
PHARMAFAIR	100,000 UNITS/GM;0.1%	A062656 001	Jul 30, 1986

NYSTATIN-TRIAMCINOLONE ACETONIDE

PHARMADERM	100,000 UNITS/GM;0.1%	A062603 001	Oct 09, 1985
------------	-----------------------	-------------	--------------

OBETICHOLIC ACID

TABLET; ORAL

OBETICHOLIC ACID

APOTEX	5MG	A214862 001	May 30, 2023
	10MG	A214862 002	May 30, 2023
LUPIN LTD	5MG	A214980 001	May 30, 2023
	10MG	A214980 002	May 30, 2023
MSN	5MG	A215017 001	May 30, 2023
	10MG	A215017 002	May 30, 2023

OCTREOTIDE ACETATE

INJECTABLE; INJECTION

OCTREOTIDE ACETATE

SHUANGCHENG	EQ 0.1MG BASE/ML	A217860 001	May 02, 2024
SUN PHARM INDS	EQ 0.05MG BASE/ML	A077329 001	Mar 04, 2008
	EQ 0.05MG BASE/ML	A077372 001	Aug 14, 2007
	EQ 0.1MG BASE/ML	A077329 002	Mar 04, 2008
	EQ 0.1MG BASE/ML	A077372 002	Aug 14, 2007
	EQ 0.2MG BASE/ML	A077330 001	Mar 04, 2008
	EQ 0.2MG BASE/ML	A077373 001	Aug 14, 2007
	EQ 0.5MG BASE/ML	A077329 003	Mar 04, 2008
	EQ 0.5MG BASE/ML	A077372 003	Aug 14, 2007
	EQ 1MG BASE/ML	A077331 001	Mar 04, 2008
	EQ 1MG BASE/ML	A077373 002	Aug 14, 2007
WOCKHARDT USA	EQ 0.2MG BASE/ML	A090986 001	May 11, 2011

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

OCTREOTIDE ACETATE

INJECTABLE; INJECTION

OCTREOTIDE ACETATE

	EQ 1MG BASE/ML	A090986 002	May 11, 2011
OCTREOTIDE ACETATE PRESERVATIVE FREE			
WOCKHARDT USA	EQ 0.05MG BASE/ML	A090985 001	May 11, 2011
	EQ 0.1MG BASE/ML	A090985 002	May 11, 2011
	EQ 0.5MG BASE/ML	A090985 003	May 11, 2011
SANDOSTATIN			
+ NOVARTIS	EQ 0.2MG BASE/ML **	N019667 004	Jun 12, 1991
+	EQ 1MG BASE/ML **	N019667 005	Jun 12, 1991

OFLOXACIN

INJECTABLE; INJECTION

FLOXIN

ORTHO MCNEIL PHARM	20MG/ML	N020087 002	Mar 31, 1992
	40MG/ML	N020087 003	Mar 31, 1992
FLOXIN IN DEXTROSE 5%			
ORTHO MCNEIL PHARM	400MG/100ML	N020087 001	Mar 31, 1992
FLOXIN IN DEXTROSE 5% IN PLASTIC CONTAINER			
ORTHO MCNEIL PHARM	4MG/ML	N020087 004	Mar 31, 1992
	400MG/100ML	N020087 005	Mar 31, 1992
OFLOXACIN			
BEDFORD	40MG/ML	A075762 001	Jan 16, 2002
SOLUTION/DROPS; OPHTHALMIC			
OFLOXACIN			
ALVOGEN	0.3%	A076830 001	Aug 31, 2004
EPIC PHARMA LLC	0.3%	A076615 001	May 14, 2004
SANDOZ	0.3%	A076231 001	May 14, 2004
	0.3%	A076848 001	Nov 25, 2008
SOLUTION/DROPS; OTIC			
FLOXIN OTIC			
+ DAIICHI	0.3% **	N020799 001	Dec 16, 1997
OFLOXACIN			
ALVOGEN	0.3%	A090395 001	Aug 11, 2009
FDC LTD	0.3%	A215038 001	Jan 19, 2022
HIKMA	0.3%	A076616 001	Mar 17, 2008
SANDOZ	0.3%	A078222 001	Mar 17, 2008
TABLET; ORAL			
FLOXIN			
+ JANSSEN PHARMS	200MG **	N019735 001	Dec 28, 1990
+	300MG **	N019735 002	Dec 28, 1990
+	400MG **	N019735 003	Dec 28, 1990
OFLOXACIN			
CHARTWELL RX	200MG	A076093 001	Sep 02, 2003
	300MG	A076093 002	Sep 02, 2003
RANBAXY LABS LTD	200MG	A076220 001	Sep 02, 2003
	300MG	A076220 002	Sep 02, 2003
	400MG	A076220 003	Sep 02, 2003

OLANZAPINE

TABLET; ORAL

OLANZAPINE

AJANTA PHARMA LTD	2.5MG	A206711 001	Aug 30, 2016
	5MG	A206711 002	Aug 30, 2016
	7.5MG	A206711 003	Aug 30, 2016
	10MG	A206711 004	Aug 30, 2016
	15MG	A206711 005	Aug 30, 2016
	20MG	A206711 006	Aug 30, 2016
HIKMA	2.5MG	A204866 001	Jun 16, 2017
	5MG	A204866 002	Jun 16, 2017
	7.5MG	A204866 003	Jun 16, 2017
	10MG	A204866 004	Jun 16, 2017
	15MG	A204866 005	Jun 16, 2017
	20MG	A204866 006	Jun 16, 2017
HISUN PHARM HANGZHOU	2.5MG	A206924 001	Dec 31, 2020
	5MG	A206924 002	Dec 31, 2020
	7.5MG	A206924 003	Dec 31, 2020
	10MG	A206924 004	Dec 31, 2020
	15MG	A206924 005	Dec 31, 2020

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

OLANZAPINETABLET; ORAL
OLANZAPINE

	20MG	A206924 006	Dec 31, 2020
INDOCO	2.5MG	A206155 001	Jul 31, 2020
	5MG	A206155 002	Jul 31, 2020
	7.5MG	A206155 003	Jul 31, 2020
	10MG	A206155 004	Jul 31, 2020
	15MG	A206155 005	Jul 31, 2020
	20MG	A206155 006	Jul 31, 2020
IVAX PHARMS INC	20MG	A077301 001	Apr 29, 2015
JIANGSU HANSOH PHARM	2.5MG	A209399 001	Sep 24, 2018
	5MG	A209399 002	Sep 24, 2018
	10MG	A209399 003	Sep 24, 2018
NATCO PHARMA	2.5MG	A076866 001	Apr 23, 2012
	5MG	A076866 002	Apr 23, 2012
	7.5MG	A076866 003	Apr 23, 2012
	10MG	A076866 004	Apr 23, 2012
	15MG	A076866 005	Apr 23, 2012
	20MG	A076866 006	Apr 23, 2012
SUN PHARM INDS	2.5MG	A091038 001	Apr 23, 2012
	5MG	A091038 002	Apr 23, 2012
	7.5MG	A091038 003	Apr 23, 2012
	10MG	A091038 004	Apr 23, 2012
	15MG	A091038 005	Apr 23, 2012
	20MG	A091038 006	Apr 23, 2012
SUNSHINE	2.5MG	A206238 001	Nov 19, 2018
	5MG	A206238 002	Nov 19, 2018
	7.5MG	A206238 003	Nov 19, 2018
	10MG	A206238 004	Nov 19, 2018
	15MG	A206238 005	Nov 19, 2018
	20MG	A206238 006	Nov 19, 2018
TEVA PHARMS	2.5MG	A076000 001	Oct 24, 2011
	5MG	A076000 002	Oct 24, 2011
	7.5MG	A076000 003	Oct 24, 2011
	10MG	A076000 004	Oct 24, 2011
	15MG	A076000 005	Oct 24, 2011
TORRENT PHARMS LTD	2.5MG	A091434 001	Apr 23, 2012
	5MG	A091434 002	Apr 23, 2012
	7.5MG	A091434 003	Apr 23, 2012
	10MG	A091434 004	Apr 23, 2012
	15MG	A091434 005	Apr 23, 2012
	20MG	A091434 006	Apr 23, 2012
TABLET, ORALLY DISINTEGRATING; ORAL			
OLANZAPINE			
AJANTA PHARMA LTD	5MG	A204320 001	May 30, 2017
	10MG	A204320 002	May 30, 2017
	15MG	A204320 003	May 30, 2017
	20MG	A204320 004	May 30, 2017
SUN PHARM INDS	5MG	A090881 001	Feb 28, 2012
	10MG	A090881 002	Feb 28, 2012
	15MG	A090881 003	Feb 28, 2012
	20MG	A090881 004	Feb 28, 2012
ZYDUS PHARMS	5MG	A202889 001	Mar 09, 2023
	10MG	A202889 002	Mar 09, 2023
	15MG	A202889 003	Mar 09, 2023
	20MG	A202889 004	Mar 09, 2023

OLAPARIB

CAPSULE; ORAL

LYNPARZA

+ ASTRAZENECA

50MG

N206162 001 Dec 19, 2014

OLICERIDINE

SOLUTION; INTRAVENOUS

OLINVIK

+ TREVENA

30MG/30ML (1MG/ML)

N210730 003 Oct 30, 2020

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

OLMESARTAN MEDOXOMIL

TABLET; ORAL

OLMESARTAN MEDOXOMIL

AMNEAL	5MG	A207480 001	Jul 11, 2023
	20MG	A207480 002	Jul 11, 2023
	40MG	A207480 003	Jul 11, 2023
JUBILANT GENERICS	5MG	A205482 001	Apr 24, 2017
	20MG	A205482 002	Apr 24, 2017
	40MG	A205482 003	Apr 24, 2017
LUPIN LTD	5MG	A206631 001	Apr 27, 2017
	20MG	A206631 002	Apr 27, 2017
	40MG	A206631 003	Apr 27, 2017
RISING	5MG	A078276 001	Oct 26, 2016
	20MG	A078276 002	Oct 26, 2016
	40MG	A078276 003	Oct 26, 2016
SANDOZ	5MG	A090237 001	Apr 13, 2020
	20MG	A090237 002	Apr 13, 2020
	40MG	A090237 003	Apr 13, 2020
TEVA PHARMS USA	5MG	A091079 001	Apr 24, 2017
	20MG	A091079 002	Apr 24, 2017
	40MG	A091079 003	Apr 24, 2017
TORRENT	5MG	A202375 001	Apr 24, 2017
	20MG	A202375 002	Apr 24, 2017
	40MG	A202375 003	Apr 24, 2017

OLOPATADINE HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

OLOPATADINE HYDROCHLORIDE

BAUSCH AND LOMB INC	EQ 0.1% BASE	A206046 001	Jul 26, 2017
	EQ 0.2% BASE	A206087 001	Dec 05, 2017
FDC LTD	EQ 0.1% BASE	A209282 001	Sep 26, 2019
FLORIDA	EQ 0.7% BASE	A208637 001	Feb 19, 2020
RISING	EQ 0.1% BASE	A204392 001	Mar 21, 2018
	EQ 0.2% BASE	A204620 001	Jun 16, 2020
SCIEGEN PHARMS INC	EQ 0.1% BASE	A204532 001	Jan 10, 2017
	EQ 0.2% BASE	A204723 001	Dec 05, 2017
ZAMBON SPA	EQ 0.1% BASE	A204706 001	Dec 07, 2015

SPRAY, METERED; NASAL

OLOPATADINE HYDROCHLORIDE

HIKMA	0.665MG/SPRAY	A213757 001	Aug 19, 2020
PATANASE			
+ NOVARTIS	0.665MG/SPRAY **	N021861 001	Apr 15, 2008

OMACETAXINE MEPESUCCINATE

POWDER; SUBCUTANEOUS

SYNRIBO

+ TEVA PHARMS INTL	3.5MG/VIAL	N203585 001	Oct 26, 2012
--------------------	------------	-------------	--------------

OMBITASVIR; PARITAPREVR; RITONAVIR

TABLET; ORAL

TECHNIVIE

+ ABBVIE	12.5MG; 75MG; 50MG **	N207931 001	Jul 24, 2015
----------	-----------------------	-------------	--------------

OMEGA-3-ACID ETHYL ESTERS

CAPSULE; ORAL

OMEGA-3-ACID ETHYL ESTERS

STRIDES SOFTGELS	1GM CONTAINS AT LEAST 900MG OF THE ETHYL ESTERS OF OMEGA-3 FATTY ACIDS	A091018 001	Jun 24, 2014
ZYDUS LIFESCIENCES	1GM CONTAINS AT LEAST 900MG OF THE ETHYL ESTERS OF OMEGA-3 FATTY ACIDS	A210107 001	Jun 14, 2019

OMEGA-3-ACID ETHYL ESTERS TYPE A

CAPSULE; ORAL

OMTRYG

+ OSMOTICA PHARM US	1.2GM CONTAINS AT LEAST 900MG OF THE ETHYL ESTERS OF OMEGA-3 FATTY ACIDS	N204977 001	Apr 23, 2014
---------------------	--	-------------	--------------

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

OMEGA-3-CARBOXYLIC ACIDS

CAPSULE;ORAL

EPANOVA

+	ASTRAZENECA	1GM CONTAINS AT LEAST 850MG OF POLYUNSATURATED FATTY ACIDS	N205060 001	May 05, 2014
---	-------------	---	-------------	--------------

OMEPRAZOLE

CAPSULE, DELAYED REL PELLETS;ORAL

OMEPRAZOLE

LUPIN LTD	40MG	A202384 001	Aug 25, 2015
NORVIUM BIOSCIENCE	10MG	A205070 001	Jun 29, 2018
	20MG	A205070 002	Jun 29, 2018
	40MG	A205070 003	Jun 29, 2018
STRIDES PHARMA	10MG	A075876 001	May 29, 2003
	20MG	A075876 002	May 29, 2003
	40MG	A075876 003	Jan 21, 2009
TEVA PHARMS USA	40MG	A204661 002	Jun 13, 2017
PRILOSEC			
+	ASTRAZENECA	10MG **	N019810 003 Oct 05, 1995
+		20MG **	N019810 001 Sep 14, 1989
+		40MG **	N019810 002 Jan 15, 1998

OMEPRAZOLE MAGNESIUM

CAPSULE, DELAYED RELEASE;ORAL

OMEPRAZOLE MAGNESIUM

AUROBINDO PHARMA LTD	EQ 20MG BASE	A213201 001	Apr 28, 2023
SPIL	EQ 20MG BASE	A210593 001	Jul 20, 2018

TABLET, DELAYED RELEASE;ORAL

OMEPRAZOLE MAGNESIUM

PERRIGO R AND D	EQ 20MG BASE	A204152 001	Jul 30, 2015
-----------------	--------------	-------------	--------------

OMEPRAZOLE; SODIUM BICARBONATE

CAPSULE;ORAL

OMEPRAZOLE AND SODIUM BICARBONATE

CHARTWELL RX	20MG;1.1GM	A204137 001	Jul 15, 2016
STRIDES PHARMA	20MG;1.1GM	A078966 001	May 25, 2010
	20MG;1.1GM	A201946 001	Jul 15, 2016
	40MG;1.1GM	A078966 002	May 25, 2010

ZEGERID

+	SALIX	20MG;1.1GM	N021849 001 Feb 27, 2006
+		40MG;1.1GM	N021849 002 Feb 27, 2006

FOR SUSPENSION;ORAL

ZEGERID

+	SALIX	20MG/PACKET;1.68GM/PACKET	N021636 001 Jun 15, 2004
+		40MG/PACKET;1.68GM/PACKET	N021636 002 Dec 21, 2004

OMIDENEPAG ISOPROPYL

SOLUTION;OPHTHALMIC

OMLONTI

+	OCUVEX THERAP	0.002%	N215092 001 Sep 22, 2022
---	---------------	--------	--------------------------

ONDANSETRON

FILM;ORAL

ZUPLENZ

+	AQUESTIVE	4MG	N022524 001 Jul 02, 2010
+		8MG	N022524 002 Jul 02, 2010

TABLET, ORALLY DISINTEGRATING;ORAL

ONDANSETRON

BARR	4MG	A076693 001	Jun 25, 2007
	8MG	A076693 002	Jun 25, 2007
IPCA LABS LTD	4MG	A209389 001	Oct 30, 2023
	8MG	A209389 002	Oct 30, 2023
MYLAN	4MG	A078139 001	Jun 25, 2007
	8MG	A078139 002	Jun 25, 2007
NESHER PHARMS	4MG	A077717 001	Jun 25, 2007
	8MG	A077717 002	Jun 25, 2007
SUN PHARM INDS LTD	4MG	A078602 001	Feb 24, 2011
	8MG	A078602 002	Feb 24, 2011
TEVA	4MG	A076810 001	Jun 25, 2007
	8MG	A076810 002	Jun 25, 2007

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ONDANSETRON

TABLET, ORALLY DISINTEGRATING;ORAL

ZOFRAN ODT

+	SANDOZ	4MG **	N020781 001	Jan 27, 1999
+		8MG **	N020781 002	Jan 27, 1999

ONDANSETRON HYDROCHLORIDE

INJECTABLE; INJECTION

ONDANSETRON HYDROCHLORIDE

AM REGENT	EQ 2MG BASE/ML	A077582 001	Dec 26, 2006
APOTEX INC	EQ 2MG BASE/ML	A077368 001	Dec 26, 2006
AVET LIFESCIENCES	EQ 2MG BASE/ML	A090424 001	Apr 16, 2010
BAXTER HLTHCARE CORP	EQ 2MG BASE/ML	A078288 001	Feb 22, 2013
CHARTWELL MOLECULAR	EQ 2MG BASE/ML	A090116 001	Apr 14, 2010
EUGIA PHARMA	EQ 2MG BASE/ML	A202599 001	Dec 21, 2012
HOSPIRA	EQ 2MG BASE/ML	A076695 001	Dec 26, 2006
	EQ 2MG BASE/ML	A077840 001	Jan 19, 2007
LANNETT CO INC	EQ 2MG BASE/ML	A090883 001	Aug 05, 2010
LUITPOLD	EQ 2MG BASE/ML	A079039 001	Nov 18, 2008
PLIVA HRVATSKA DOO	EQ 2MG BASE/ML	A077544 001	Dec 26, 2006
RISING	EQ 2MG BASE/ML	A204906 001	Jul 31, 2017
SAGENT PHARMS	EQ 2MG BASE/ML	A078180 001	Mar 26, 2007
STERISCIENCE SPECLTS	EQ 2MG BASE/ML	A078257 001	Apr 23, 2008
SUN PHARM INDS (IN)	EQ 2MG BASE/ML	A077172 001	Dec 26, 2006
TEVA	EQ 2MG BASE/ML	A076876 001	Nov 22, 2006

ONDANSETRON HYDROCHLORIDE AND DEXTROSE IN PLASTIC CONTAINER

HOSPIRA	EQ 0.64MG BASE/ML	A076978 001	Feb 26, 2007
---------	-------------------	-------------	--------------

ONDANSETRON HYDROCHLORIDE PRESERVATIVE FREE

ACCORD HLTHCARE	EQ 2MG BASE/ML	A206845 001	Mar 10, 2016
AM REGENT	EQ 2MG BASE/ML	A077387 001	Dec 26, 2006
	EQ 2MG BASE/ML	A079032 001	Nov 18, 2008
APOTEX INC	EQ 2MG BASE/ML	A077343 001	Dec 26, 2006
AVET LIFESCIENCES	EQ 2MG BASE/ML	A078945 001	Jan 03, 2013
EUGIA PHARMA	EQ 2MG BASE/ML	A202600 001	Dec 21, 2012
HIKMA FARMACEUTICA	EQ 2MG BASE/ML	A076780 001	Dec 26, 2006
HOSPIRA	EQ 2MG BASE/ML	A076696 001	Dec 26, 2006
STERISCIENCE SPECLTS	EQ 2MG BASE/ML	A078244 001	Apr 23, 2008
SUN PHARM INDS LTD	EQ 2MG BASE/ML	A077173 001	Dec 26, 2006
TARO PHARMS IRELAND	EQ 2MG BASE/ML	A078014 001	Mar 21, 2008
TEVA	EQ 2MG BASE/ML	A076759 001	Nov 22, 2006

ZOFRAN

+	SANDOZ	EQ 2MG BASE/ML **	N020007 001	Jan 04, 1991
---	--------	-------------------	-------------	--------------

ZOFRAN AND DEXTROSE IN PLASTIC CONTAINER

+	GLAXOSMITHKLINE	EQ 0.64MG BASE/ML **	N020403 001	Jan 31, 1995
---	-----------------	----------------------	-------------	--------------

ZOFRAN PRESERVATIVE FREE

+	SANDOZ	EQ 2MG BASE/ML **	N020007 003	Dec 10, 1993
---	--------	-------------------	-------------	--------------

SOLUTION;ORAL

ONDANSETRON HYDROCHLORIDE

PHARM ASSOC	EQ 4MG BASE/5ML	A078127 001	Jun 25, 2007
TARO	EQ 4MG BASE/5ML	A077009 001	Nov 30, 2007

ZOFRAN

+	SANDOZ	EQ 4MG BASE/5ML	N020605 001	Jan 24, 1997
---	--------	-----------------	-------------	--------------

TABLET;ORAL

ONDANSETRON HYDROCHLORIDE

AUROBINDO PHARMA	EQ 24MG BASE	A078539 003	Jul 31, 2007
CHARTWELL RX	EQ 4MG BASE	A077517 001	Jun 25, 2007
	EQ 8MG BASE	A077517 002	Jun 25, 2007
	EQ 24MG BASE	A077517 003	Jun 25, 2007
DR REDDYS LABS LTD	EQ 16MG BASE	A076183 004	Dec 26, 2006
	EQ 24MG BASE	A076183 001	Dec 26, 2006
GLENMARK PHARMS LTD	EQ 24MG BASE	A077535 003	Jun 25, 2007
HIKMA INTL PHARMS	EQ 4MG BASE	A077545 001	Sep 06, 2007
	EQ 8MG BASE	A077545 002	Sep 06, 2007
	EQ 24MG BASE	A077545 003	Sep 06, 2007
IPCA LABS LTD	EQ 4MG BASE	A203761 001	Jan 23, 2014
	EQ 8MG BASE	A203761 002	Jan 23, 2014
PLIVA HRVATSKA DOO	EQ 4MG BASE	A077112 001	Jun 25, 2007
	EQ 8MG BASE	A077112 002	Jun 25, 2007
	EQ 24MG BASE	A077112 003	Jun 25, 2007
RISING	EQ 4MG BASE	A076930 001	Jun 25, 2007

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ONDANSETRON HYDROCHLORIDE

TABLET; ORAL

ONDANSETRON HYDROCHLORIDE

	EQ 8MG BASE	A076930 002	Jun 25, 2007
	EQ 24MG BASE	A076930 004	Jun 25, 2007
SUN PHARM INDS (IN)	EQ 4MG BASE	A077050 001	Jun 25, 2007
	EQ 8MG BASE	A077050 002	Jun 25, 2007
TARO	EQ 4MG BASE	A077729 001	Mar 28, 2011
	EQ 8MG BASE	A077729 002	Mar 28, 2011
	EQ 24MG BASE	A077729 003	Mar 28, 2011
TEVA	EQ 4MG BASE	A076252 001	Jun 25, 2007
	EQ 8MG BASE	A076252 002	Jun 25, 2007
	EQ 24MG BASE	A076252 003	Jun 25, 2007
ZOFRAN			
+ SANDOZ	EQ 4MG BASE **	N020103 001	Dec 31, 1992
+	EQ 8MG BASE **	N020103 002	Dec 31, 1992
+	EQ 24MG BASE **	N020103 003	Aug 27, 1999

ORPHENADRINE CITRATE

INJECTABLE; INJECTION

NORFLEX

+ PAI HOLDINGS PHARM	30MG/ML **	N013055 001	
ORPHENADRINE CITRATE			
WATSON LABS	30MG/ML	A087062 001	
TABLET, EXTENDED RELEASE; ORAL			
NORFLEX			
+ BAUSCH	100MG **	N012157 001	
ORPHENADRINE CITRATE			
ASCOT	100MG	A088067 001	Apr 06, 1983
SANDOZ	100MG	A085046 001	
STEVENS J	100MG	A040368 001	Jun 23, 2000
WATSON LABS	100MG	A084303 001	

ORPHENADRINE HYDROCHLORIDE

TABLET; ORAL

DISIPAL

3M 50MG N010653 001

OSELTAMIVIR PHOSPHATE

CAPSULE; ORAL

OSELTAMIVIR PHOSPHATE

ACCORD HLTHCARE	EQ 30MG BASE	A214726 001	Jul 25, 2022
	EQ 45MG BASE	A214726 002	Jul 25, 2022
	EQ 75MG BASE	A214726 003	Jul 25, 2022
RISING	EQ 30MG BASE	A210157 001	Jan 21, 2021
	EQ 45MG BASE	A210157 002	Jan 21, 2021
	EQ 75MG BASE	A210157 003	Jan 21, 2021

FOR SUSPENSION; ORAL

OSELTAMIVIR PHOSPHATE

LEADING	EQ 6MG BASE/ML	A214949 001	Feb 28, 2022
TAMIFLU			
ROCHE	EQ 12MG BASE/ML	N021246 001	Dec 14, 2000

OSILODROSTAT PHOSPHATE

TABLET; ORAL

ISTURISA

+ RECORDATI RARE EQ 10MG BASE N212801 003 Mar 06, 2020

OXACILLIN SODIUM

CAPSULE; ORAL

BACTOCILL

GLAXOSMITHKLINE	EQ 250MG BASE	A061336 001	
	EQ 250MG BASE	A062241 001	
	EQ 500MG BASE	A061336 002	
	EQ 500MG BASE	A062241 002	

OXACILLIN SODIUM

ANI PHARMS	EQ 250MG BASE	A062222 001	
	EQ 500MG BASE	A062222 002	
APOTHECON	EQ 250MG BASE	A061450 002	
	EQ 500MG BASE	A061450 001	

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

OXACILLIN SODIUM

CAPSULE; ORAL

PROSTAPHLIN

APOTHECON EQ 500MG BASE N050118 002

FOR SOLUTION; ORAL

BACTOCILL

GLAXOSMITHKLINE EQ 250MG BASE/5ML A062321 001

OXACILLIN SODIUM

APOTHECON EQ 250MG BASE/5ML A061457 001

TEVA EQ 250MG BASE/5ML A062252 001

PROSTAPHLIN

APOTHECON EQ 250MG BASE/5ML N050194 001

INJECTABLE; INJECTION

BACTOCILL

GLAXOSMITHKLINE EQ 500MG BASE/VIAL ** A061334 009 Mar 26, 1982

EQ 1GM BASE/VIAL ** A061334 006 Mar 26, 1982

EQ 1GM BASE/VIAL ** A062736 001 Dec 19, 1986

EQ 2GM BASE/VIAL ** A061334 007 Mar 26, 1982

EQ 2GM BASE/VIAL ** A062736 002 Dec 19, 1986

EQ 4GM BASE/VIAL ** A061334 008 Mar 26, 1982

EQ 10GM BASE/VIAL ** A061334 010

OXACILLIN SODIUM

+ APOTHECON EQ 250MG BASE/VIAL ** N050195 001

+ EQ 500MG BASE/VIAL ** N050195 002

+ EQ 1GM BASE/VIAL ** N050195 003

+ EQ 2GM BASE/VIAL ** N050195 004

+ EQ 4GM BASE/VIAL ** N050195 005

ELKINS SINN EQ 250MG BASE/VIAL A062711 001 Feb 03, 1989

EQ 500MG BASE/VIAL A062711 002 Feb 03, 1989

EQ 1GM BASE/VIAL A062711 003 Feb 03, 1989

EQ 2GM BASE/VIAL A062711 004 Feb 03, 1989

EQ 4GM BASE/VIAL A062711 005 Feb 03, 1989

EQ 10GM BASE/VIAL A062711 006 Feb 03, 1989

HOSPIRA EQ 1GM BASE/VIAL A203950 001 Dec 11, 2015

EQ 2GM BASE/VIAL A203950 002 Dec 11, 2015

ISTITUTO BIO ITA SPA EQ 125MG BASE/VIAL A062798 003 Dec 11, 1995

EQ 250MG BASE/VIAL A062798 004 Dec 11, 1995

EQ 500MG BASE/VIAL A062798 005 Dec 11, 1995

EQ 1GM BASE/VIAL A062798 001 Dec 11, 1995

EQ 2GM BASE/VIAL A062798 002 Dec 11, 1995

PIRAMAL CRITICAL EQ 1GM BASE/VIAL A206681 001 Sep 11, 2017

EQ 2GM BASE/VIAL A206681 002 Sep 11, 2017

EQ 10GM BASE/VIAL A206760 001 Oct 26, 2017

SAGENT PHARMS EQ 1GM BASE/VIAL A091246 001 Mar 30, 2012

EQ 2GM BASE/VIAL A091246 002 Mar 30, 2012

SANDOZ EQ 250MG BASE/VIAL A061490 001

EQ 500MG BASE/VIAL A061490 002

EQ 1GM BASE/VIAL A061490 003

EQ 2GM BASE/VIAL A061490 004

EQ 10GM BASE/VIAL A061490 006 May 09, 1991

STERISCIENCE SPECLTS EQ 1GM BASE/VIAL A091486 001 Aug 25, 2014

WATSON LABS INC EQ 250MG BASE/VIAL A062856 001 Oct 26, 1988

EQ 500MG BASE/VIAL A062856 002 Oct 26, 1988

EQ 1GM BASE/VIAL A062856 003 Oct 26, 1988

EQ 2GM BASE/VIAL A062856 004 Oct 26, 1988

EQ 4GM BASE/VIAL A062856 005 Oct 26, 1988

EQ 10GM BASE/VIAL A062984 001 Sep 29, 1988

POWDER; INTRAVENOUS

OXACILLIN SODIUM

SANDOZ EQ 1GM BASE/VIAL A062737 001 Dec 23, 1986

EQ 2GM BASE/VIAL A062737 002 Dec 23, 1986

OXALIPLATIN

INJECTABLE; INTRAVENOUS

ELOXATIN

+ SANOFI AVENTIS US 50MG/VIAL ** N021492 001 Aug 09, 2002

+ 50MG/10ML (5MG/ML) N021759 001 Jan 31, 2005

+ 100MG/VIAL ** N021492 002 Aug 09, 2002

+ 100MG/20ML (5MG/ML) N021759 002 Jan 31, 2005

+ 200MG/40ML (5MG/ML) ** N021759 003 Nov 17, 2006

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

OXALIPLATIN

INJECTABLE; INTRAVENOUS

OXALIPLATIN

ACCORD HLTHCARE	200MG/40ML (5MG/ML)	A207474	003	Mar 21, 2017
ACTAVIS TOTOWA	50MG/VIAL	A078803	001	Aug 08, 2012
	100MG/VIAL	A078803	002	Aug 08, 2012
AM REGENT	50MG/10ML (5MG/ML)	A204378	001	May 12, 2017
	100MG/20ML (5MG/ML)	A204378	002	May 12, 2017
CHARTWELL MOLECULAR	50MG/10ML (5MG/ML)	A208523	001	Feb 10, 2017
	100MG/20ML (5MG/ML)	A208523	002	Feb 10, 2017
EUGIA PHARMA	50MG/10ML (5MG/ML)	A205529	001	Sep 06, 2017
	100MG/20ML (5MG/ML)	A205529	002	Sep 06, 2017
FRESENIUS KABI ONCOL	50MG/VIAL	A078810	001	Aug 07, 2009
	100MG/VIAL	A078810	002	Aug 07, 2009
FRESENIUS KABI USA	200MG/40ML (5MG/ML)	A090030	003	Jan 31, 2017
GLAND	200MG/40ML (5MG/ML)	A207325	003	Oct 18, 2017
HOSPIRA INC	50MG/VIAL	A078815	001	Sep 30, 2009
	100MG/VIAL	A078815	002	Sep 30, 2009
MYLAN LABS LTD	200MG/40ML (5MG/ML)	A091358	003	Nov 14, 2017
NORVIUM BIOSCIENCE	50MG/VIAL	A200979	001	Aug 08, 2012
	100MG/VIAL	A200979	002	Aug 08, 2012
SANDOZ	50MG/10ML (5MG/ML)	A078812	001	Aug 07, 2009
	50MG/VIAL	A090849	001	Apr 28, 2011
	100MG/20ML (5MG/ML)	A078812	002	Aug 07, 2009
	100MG/VIAL	A090849	002	Apr 28, 2011
SUN PHARM	50MG/VIAL	A078818	001	Aug 07, 2009
	50MG/10ML (5MG/ML)	A202922	001	Apr 08, 2014
	100MG/VIAL	A078818	002	Aug 07, 2009
	100MG/20ML (5MG/ML)	A202922	002	Apr 08, 2014
	200MG/40ML (5MG/ML)	A202922	003	Feb 15, 2019

OXAMNIQUINE

CAPSULE; ORAL

VANSIL

PFIZER	250MG	N018069	001	
--------	-------	---------	-----	--

OXANDROLONE

TABLET; ORAL

OXANDROLONE

PAR PHARM	2.5MG	A077827	001	Jun 22, 2007
	10MG	A077827	002	Jun 22, 2007
ROXANE	2.5MG	A077249	001	Jul 10, 2007
	10MG	A077249	002	Jul 10, 2007
SANDOZ	2.5MG	A076897	001	Dec 01, 2006
	10MG	A076897	002	Dec 01, 2006
UPSHER SMITH LABS	2.5MG	A076761	001	Dec 01, 2006
	10MG	A078033	001	Mar 22, 2007

OXAPROZIN

TABLET; ORAL

OXAPROZIN

ACTAVIS ELIZABETH	600MG	A075843	001	Oct 03, 2001
BEXIMCO PHARMS USA	600MG	A075842	001	Apr 12, 2001
IVAX SUB TEVA PHARMS	600MG	A075846	001	May 13, 2002
MYLAN PHARMS INC	600MG	A075847	001	Feb 28, 2001
NORVIUM BIOSCIENCE	600MG	A075851	001	Aug 17, 2001
SANDOZ	600MG	A075850	001	Apr 27, 2001
SUN PHARM INDS INC	600MG	A075844	001	Jan 03, 2002
TEVA	600MG	A075849	001	Jul 03, 2002
WATSON LABS	600MG	A075848	001	Feb 09, 2001

OXAPROZIN POTASSIUM

TABLET; ORAL

DAYPRO ALTA

PFIZER	600MG	N020776	001	Oct 17, 2002
--------	-------	---------	-----	--------------

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

OXAZEPAM

CAPSULE; ORAL

OXAZEPAM

AM THERAP	10MG	A071955 001	Mar 03, 1988
	15MG	A071956 001	Mar 03, 1988
	30MG	A071957 001	Mar 03, 1988
IVAX SUB TEVA PHARMS	10MG	A070943 001	Aug 03, 1987
	15MG	A070944 001	Aug 03, 1987
	30MG	A070945 001	Aug 03, 1987
MYLAN	10MG	A071713 001	Oct 20, 1987
	15MG	A071714 001	Oct 20, 1987
	30MG	A071715 001	Oct 20, 1987
WATSON LABS	15MG	A072953 001	Sep 28, 1990
	30MG	A072954 001	Sep 28, 1990
WATSON LABS TEVA	10MG	A072952 001	Sep 28, 1990
SERAX			
+ ALPHARMA US PHARMS	10MG **	N015539 002	
+	15MG **	N015539 004	
+	30MG **	N015539 006	

ZAXOPAM

QUANTUM PHARMICS	10MG	A070650 001	Mar 01, 1988
	15MG	A070640 001	Mar 01, 1988
	30MG	A070641 001	Mar 01, 1988

TABLET; ORAL

OXAZEPAM

PARKE DAVIS	15MG	A071508 001	Feb 02, 1987
SUN PHARM INDUSTRIES	15MG	A070683 001	Jan 16, 1987
WATSON LABS	15MG	A071494 001	Apr 21, 1987

SERAX

ALPHARMA US PHARMS	15MG **	N015539 008	
--------------------	---------	-------------	--

OXCARBAZEPINE

SUSPENSION; ORAL

OXCARBAZEPINE

HIKMA	300MG/5ML	A201193 001	Oct 03, 2012
RENEW PHARMS	300MG/5ML	A211420 001	Jul 09, 2021

TABLET; ORAL

OXCARBAZEPINE

HIKMA	150MG	A077795 001	Oct 09, 2007
	300MG	A077795 002	Oct 09, 2007
	600MG	A077795 003	Oct 09, 2007
JUBILANT CADISTA	150MG	A090239 001	Jan 25, 2010
	300MG	A090239 002	Jan 25, 2010
	600MG	A090239 003	Jan 25, 2010
ZYDUS	150MG	A211747 001	Jul 03, 2023
	300MG	A211747 002	Jul 03, 2023
	600MG	A211747 003	Jul 03, 2023

TABLET, EXTENDED RELEASE; ORAL

OXCARBAZEPINE

AJANTA PHARMA LTD	150MG	A217659 001	Feb 22, 2024
	300MG	A217659 002	Feb 22, 2024
	600MG	A217659 003	Feb 22, 2024
RICONPHARMA LLC	150MG	A215796 001	Nov 22, 2024
	300MG	A215796 002	Nov 22, 2024
	600MG	A215796 003	Nov 22, 2024

OPRENOLOL HYDROCHLORIDE

CAPSULE; ORAL

TRASICOR

NOVARTIS	20MG	N018166 001	Dec 28, 1983
	40MG	N018166 002	Dec 28, 1983
	80MG	N018166 003	Dec 28, 1983
	160MG	N018166 004	Dec 28, 1983

OXTRIPHYLLINE

SOLUTION; ORAL

CHOLEDYL

PARKE DAVIS	100MG/5ML	N009268 012	Nov 27, 1984
-------------	-----------	-------------	--------------

OXTRIPHYLLINE

MORTON GROVE	100MG/5ML	A088243 001	Dec 05, 1983
--------------	-----------	-------------	--------------

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

OXTRIPHYLLINE

SYRUP;ORAL

CHOLEDYL

PARKE DAVIS 50MG/5ML

N009268 011

OXTRIPHYLLINE PEDIATRIC

MORTON GROVE 50MG/5ML

A088242 001 Dec 05, 1983

TABLET, DELAYED RELEASE;ORAL

CHOLEDYL

PARKE DAVIS 100MG
200MGN009268 003
N009268 007

OXTRIPHYLLINE

WATSON LABS 100MG
200MGA087866 001 Aug 25, 1983
A087835 001 Aug 25, 1983

TABLET, EXTENDED RELEASE;ORAL

CHOLEDYL SA

WARNER CHILCOTT LLC 400MG
600MGA087863 001 May 24, 1983
A086742 001OXYBUTYNIN

FILM, EXTENDED RELEASE;TRANSDERMAL

OXYBUTYNIN

BARR LABS DIV TEVA 3.9MG/24HR

A090526 001 Mar 04, 2014

GEL, METERED;TRANSDERMAL

GELNIQUE 3%

+ ALLERGAN 3%

N202513 001 Dec 07, 2011

OXYBUTYNIN CHLORIDE

GEL;TRANSDERMAL

GELNIQUE

+ ABBVIE 10% (100MG/PACKET) **

N022204 001 Jan 27, 2009

OXYBUTYNIN CHLORIDE

CHARTWELL RX 10% (100MG/PACKET)

A207329 001 May 31, 2018

SYRUP;ORAL

DITROPAN

+ ORTHO MCNEIL JANSSEN 5MG/5ML **

N018211 001

OXYBUTYNIN CHLORIDE

LANNETT CO INC 5MG/5ML
PHARM ASSOC 5MG/5ML
5MG/5ML
PHARMOBEDIANT CNSLTG 5MG/5MLA076682 001 Dec 28, 2004
A074997 001 Oct 15, 1997
A075137 001 Dec 18, 1998
A074868 001 Feb 12, 1997

TABLET;ORAL

DITROPAN

+ JANSSEN PHARMS 5MG **

N017577 001

OXYBUTYNIN CHLORIDE

AVET LIFESCIENCES 5MG
HIBROW HLTHCARE 5MG
QUANTUM PHARMICS 5MG
USL PHARMA 5MG
WATSON LABS 5MGA211682 001 May 10, 2019
A211062 001 Feb 06, 2019
A072296 001 Dec 08, 1988
A070746 001 Mar 10, 1988
A072485 001 Apr 19, 1989

TABLET, EXTENDED RELEASE;ORAL

DITROPAN XL

+ JANSSEN PHARMS 5MG **
+ 10MG **
+ 15MG **N020897 001 Dec 16, 1998
N020897 002 Dec 16, 1998
N020897 003 Jun 22, 1999

OXYBUTYNIN CHLORIDE

IMPAX PHARMS 5MG
10MG
15MG
NORVIUM BIOSCIENCE 5MG
10MG
15MGA076745 002 May 09, 2007
A076745 003 May 09, 2007
A076745 001 Nov 09, 2006
A076702 001 Nov 09, 2006
A076644 001 Nov 09, 2006
A076644 002 May 10, 2007OXYCODONE HYDROCHLORIDE

CAPSULE;ORAL

OXYCODONE HYDROCHLORIDE

DR REDDYS LABS SA 5MG
LANNETT CO INC 5MGA203107 001 Jul 26, 2012
A203823 001 Aug 01, 2014

SOLUTION;ORAL

OXYCODONE HYDROCHLORIDE

ANI PHARMS 100MG/5ML
AUROLIFE PHARMA LLC 5MG/5MLA203447 001 Aug 30, 2017
A212429 001 Jan 27, 2020

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

OXYCODONE HYDROCHLORIDE

SOLUTION;ORAL

OXYCODONE HYDROCHLORIDE

	100MG/5ML	A212429	002	Jan 27, 2020
CHARTWELL MOLECULAR	100MG/5ML	A204085	001	Sep 09, 2014
DR REDDYS LABS SA	100MG/5ML	A204092	001	Jun 05, 2014
HIKMA	5MG/5ML	A208817	001	Aug 10, 2017
	100MG/5ML	A203208	001	Jul 12, 2013
	100MG/5ML	A208795	001	Aug 07, 2017
NOVEL LABS INC	100MG/5ML	A204603	001	Apr 29, 2015
RHODES PHARMS	100MG/5ML	A205853	001	Apr 29, 2020
VISTAPHARM	100MG/5ML	A202537	001	Jul 30, 2012
+ VISTAPHARM LLC	5MG/5ML **	N201194	001	Jan 12, 2012
XTRTRIUM LABS INC	5MG/5ML	A206456	001	Jun 16, 2015

TABLET;ORAL

OXAYDO

+ ZYLA	5MG	N202080	001	Jun 17, 2011
+	7.5MG	N202080	002	Jun 17, 2011

OXYCODONE HYDROCHLORIDE

ACTAVIS ELIZABETH	5MG	A076636	003	Apr 07, 2015
	15MG	A076636	001	Feb 06, 2004
	30MG	A076636	002	Feb 06, 2004
DR REDDYS LABS SA	5MG	A091313	001	Feb 18, 2011
	10MG	A091313	004	Apr 29, 2016
	15MG	A091313	002	Feb 18, 2011
	20MG	A091313	005	Apr 29, 2016
	30MG	A091313	003	Feb 18, 2011
NESHER PHARMS	5MG	A077290	001	Dec 08, 2005
	10MG	A077290	002	Dec 08, 2005
	15MG	A077290	003	Dec 08, 2005
	20MG	A077290	004	Dec 08, 2005
	30MG	A077290	005	Dec 08, 2005
SUN PHARM INDS INC	5MG	A090659	001	Apr 10, 2009
	10MG	A090659	005	Nov 06, 2012
	15MG	A090659	002	Apr 10, 2009
	20MG	A090659	004	Nov 06, 2012
	30MG	A090659	003	Apr 10, 2009

TABLET, EXTENDED RELEASE;ORAL

ROXICODONE

ROXANE	10MG	N020932	001	Oct 26, 1998
	30MG	N020932	002	Oct 26, 1998

OXYMETAZOLINE HYDROCHLORIDE

CREAM;TOPICAL

OXYMETAZOLINE HYDROCHLORIDE

TARO	1%	A213584	001	Oct 04, 2021
------	----	---------	-----	--------------

SOLUTION/DROPS;OPHTHALMIC

OCUCLEAR

BAYER HEALTHCARE LLC	0.025%	N018471	001	May 30, 1986
----------------------	--------	---------	-----	--------------

OXYMETHOLONE

TABLET;ORAL

ANADROL-50

+ NORVIUM BIOSCIENCE	50MG	N016848	001	
----------------------	------	---------	-----	--

OXYMORPHONE HYDROCHLORIDE

INJECTABLE;INJECTION

OPANA

+ ENDO PHARMS	1MG/ML	N011707	002	
	1.5MG/ML	N011707	001	

SUPPOSITORY;RECTAL

NUMORPHAN

ENDO PHARMS	5MG	N011738	004	
-------------	-----	---------	-----	--

TABLET;ORAL

OPANA

+ ENDO PHARMS	5MG **	N021611	001	Jun 22, 2006
+	10MG **	N021611	002	Jun 22, 2006

OXYMORPHONE HYDROCHLORIDE

SPECGX LLC	5MG	A202321	001	Apr 25, 2013
	10MG	A202321	002	Apr 25, 2013

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

OXYMORPHONE HYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL

OPANA ER

+	ENDO OPERATIONS	5MG **	N021610 001	Jun 22, 2006
+		7.5MG **	N021610 005	Feb 29, 2008
+		10MG **	N021610 002	Jun 22, 2006
+		15MG **	N021610 006	Feb 29, 2008
+		20MG **	N021610 003	Jun 22, 2006
+		30MG **	N021610 007	Feb 29, 2008
+		40MG **	N021610 004	Jun 22, 2006
+	ENDO PHARMS	5MG	N201655 001	Dec 09, 2011
+		7.5MG	N201655 002	Dec 09, 2011
+		10MG	N201655 003	Dec 09, 2011
+		15MG	N201655 004	Dec 09, 2011
+		20MG	N201655 005	Dec 09, 2011
+		30MG	N201655 006	Dec 09, 2011
+		40MG	N201655 007	Dec 09, 2011

OXYMORPHONE HYDROCHLORIDE

ACTAVIS ELIZABETH

	5MG	A079046 003	Jul 11, 2013
	7.5MG	A079046 001	Dec 13, 2010
	10MG	A079046 004	Jul 11, 2013
	15MG	A079046 002	Dec 13, 2010
	20MG	A079046 005	Jul 11, 2013
	30MG	A079046 006	Jul 11, 2013
	40MG	A079046 007	Jul 11, 2013

HIKMA

	5MG	A200822 002	Jul 15, 2013
	7.5MG	A200822 003	Jul 15, 2013
	10MG	A200822 004	Jul 15, 2013
	15MG	A200822 005	Jul 15, 2013
	20MG	A200822 006	Jul 15, 2013
	30MG	A200822 007	Jul 15, 2013
	40MG	A200822 001	Jul 15, 2013

PAR PHARM

	5MG	A200792 001	Oct 24, 2014
	7.5MG	A200792 002	Oct 24, 2014
	10MG	A200792 003	Oct 24, 2014
	15MG	A200792 004	Oct 24, 2014
	20MG	A200792 005	Oct 24, 2014
	30MG	A200792 006	Oct 24, 2014
	40MG	A200792 007	Oct 24, 2014

SPECGX LLC

	5MG	A202946 001	Jun 27, 2014
	7.5MG	A202946 002	Jun 27, 2014
	10MG	A202946 003	Jun 27, 2014
	15MG	A202946 004	Jun 27, 2014
	20MG	A202946 005	Jun 27, 2014
	30MG	A202946 006	Jun 27, 2014
	40MG	A202946 007	Jun 27, 2014

SUN PHARM INDS LTD

	5MG	A203506 001	Apr 24, 2015
	7.5MG	A203506 002	Apr 24, 2015
	10MG	A203506 003	Apr 24, 2015
	15MG	A203506 004	Apr 24, 2015
	20MG	A203506 005	Apr 24, 2015
	30MG	A203506 006	Apr 24, 2015
	40MG	A203506 007	Apr 24, 2015

OXYPHENBUTAZONE

TABLET;ORAL

OXYPHENBUTAZONE

WATSON LABS

	100MG	A088399 001	Sep 17, 1984
--	-------	-------------	--------------

TANDEARIL

NOVARTIS

	100MG	N012542 004	Sep 03, 1982
--	-------	-------------	--------------

OXYPHENCYCLIMINE HYDROCHLORIDE

TABLET;ORAL

DARICON

PFIZER

	10MG	N011612 001	
--	------	-------------	--

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

OXYPHENONIUM BROMIDE

TABLET; ORAL

ANTRENYL

NOVARTIS

5MG

N008492 002

OXYTETRACYCLINE

TABLET; ORAL

TERRAMYCIN

PFIZER

250MG

N050287 001

OXYTETRACYCLINE CALCIUM

SYRUP; ORAL

TERRAMYCIN

PFIZER

EQ 125MG BASE/5ML

A060595 001

OXYTETRACYCLINE HYDROCHLORIDE

CAPSULE; ORAL

OXY-KESSO-TETRA

FERRANTE

EQ 250MG BASE

A060179 001

OXYTETRACYCLINE HYDROCHLORIDE

HIKMA

EQ 250MG BASE

A060770 001

IMPAX LABS

EQ 250MG BASE

A060760 001

PROTER

EQ 250MG BASE

A060869 001

PUREPAC PHARM

EQ 250MG BASE

A060634 001

TERRAMYCIN

PFIZER

EQ 125MG BASE

N050286 001

EQ 250MG BASE

N050286 002

INJECTABLE; INJECTION

TERRAMYCIN

PFIZER

EQ 250MG BASE/VIAL

A060586 001

EQ 500MG BASE/VIAL

A060586 002

OXYTETRACYCLINE HYDROCHLORIDE; POLYMYXIN B SULFATE

OINTMENT; OPHTHALMIC

TERRAMYCIN W/ POLYMYXIN B SULFATE

CASPER PHARMA LLC

EQ 5MG BASE/GM;10,000 UNITS/GM

N061015 001

OINTMENT; OTIC

TERRAMYCIN W/ POLYMYXIN

PFIZER

EQ 5MG BASE/GM;10,000 UNITS/GM

A061841 001

TABLET; VAGINAL

TERRAMYCIN-POLYMYXIN

PFIZER

EQ 100MG BASE;100,000 UNITS

A061009 001

OXYTOCIN

INJECTABLE; INJECTION

OXYTOCIN

DR REDDYS

10USP UNITS/ML (10USP UNITS/ML)

A077453 001 Jan 24, 2008

100USP UNITS/10ML (10USP UNITS/ML)

A077453 002 Jan 24, 2008

OXYTOCIN 10 USP UNITS IN DEXTROSE 5%

+ ABBOTT

1USP UNITS/100ML **

N019185 004 Mar 29, 1985

+

2USP UNITS/100ML **

N019185 003 Mar 29, 1985

OXYTOCIN 20 USP UNITS IN DEXTROSE 5%

+ ABBOTT

2USP UNITS/100ML **

N019185 002 Mar 29, 1985

OXYTOCIN 5 USP UNITS IN DEXTROSE 5%

+ ABBOTT

1USP UNITS/100ML **

N019185 001 Mar 29, 1985

SYNTOCINON

NOVARTIS

10USP UNITS/ML

N018245 001

SOLUTION; NASAL

SYNTOCINON

RTRX

40USP UNITS/ML

N012285 001

OZENOXACIN

CREAM; TOPICAL

XEPI

+ FERRER INTERNACIONAL 1%

N208945 001 Dec 11, 2017

PACLITAXEL

INJECTABLE; INJECTION

PACLITAXEL

ACCORD HLTHCARE

6MG/ML

A075436 001 Nov 12, 2004

DASH PHARMS

6MG/ML

A091540 001 Sep 29, 2011

HOSPIRA

6MG/ML

A076233 001 Aug 01, 2002

NORVIUM BIOSCIENCE

6MG/ML

A075278 001 Jan 25, 2002

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PACLITAXEL

INJECTABLE; INJECTION

PACLITAXEL

PLIVA LACHEMA	6MG/ML	A077413	001	Mar 12, 2008
SANDOZ	6MG/ML	A078167	001	Dec 26, 2007
TEVA PHARMS USA	6MG/ML	A075297	001	Jan 25, 2002

TAXOL

+ HQ SPCLT PHARMA	6MG/ML **	N020262	001	Dec 29, 1992
-------------------	-----------	---------	-----	--------------

PALBOCICLIB

TABLET; ORAL

PALBOCICLIB

SYNTHON PHARMS INC	75MG	A215570	001	Jun 05, 2024
	100MG	A215570	002	Jun 05, 2024
	125MG	A215570	003	Jun 05, 2024

PALIPERIDONE

TABLET, EXTENDED RELEASE; ORAL

INVEGA

+ JANSSEN PHARMS	1.5MG **	N021999	006	Aug 26, 2008
------------------	----------	---------	-----	--------------

+	12MG **	N021999	004	Dec 19, 2006
---	---------	---------	-----	--------------

PALIPERIDONE

ACTAVIS LABS FL INC	1.5MG	A202645	001	Aug 03, 2015
	3MG	A202645	002	Aug 03, 2015
	6MG	A202645	003	Aug 03, 2015
	9MG	A202645	004	Aug 03, 2015
AJANTA PHARMA LTD	3MG	A218514	002	Jun 26, 2024
	6MG	A218514	003	Jun 26, 2024
	9MG	A218514	004	Jun 26, 2024

PALONOSETRON HYDROCHLORIDE

CAPSULE; ORAL

ALOXI

+ HELSINN HLTHCARE	EQ 0.5MG BASE **	N022233	001	Aug 22, 2008
--------------------	------------------	---------	-----	--------------

INJECTABLE; INTRAVENOUS

ALOXI

+ HELSINN HLTHCARE	EQ 0.075MG BASE/1.5ML (EQ 0.05MG	N021372	002	Feb 29, 2008
--------------------	----------------------------------	---------	-----	--------------

	BASE/ML) **			
--	-------------	--	--	--

+	EQ 0.25MG BASE/5ML (EQ 0.05MG BASE/ML)	N021372	001	Jul 25, 2003
---	--	---------	-----	--------------

	**			
--	----	--	--	--

PALONOSETRON HYDROCHLORIDE

ACCORD HLTHCARE	EQ 0.25MG BASE/5ML (EQ 0.05MG BASE/ML)	A204615	001	Mar 15, 2021
CIPLA	EQ 0.25MG BASE/5ML (EQ 0.05MG BASE/ML)	A206396	001	Sep 19, 2018
DR REDDYS	EQ 0.075MG BASE/1.5ML (EQ 0.05MG	A201533	001	Apr 21, 2016
	BASE/ML)			
HOSPIRA	EQ 0.075MG BASE/1.5ML (EQ 0.05MG	A207005	002	Sep 19, 2018
	BASE/ML)			
NOVAST LABS	EQ 0.25MG BASE/5ML (EQ 0.05MG BASE/ML)	A208789	001	May 22, 2020
QILU PHARM HAINAN	EQ 0.075MG BASE/1.5ML (EQ 0.05MG	A205648	002	Sep 19, 2018
	BASE/ML)			
TEVA PHARMS USA	EQ 0.075MG BASE/1.5ML (EQ 0.05MG	A090713	002	Mar 23, 2018
	BASE/ML)			

SOLUTION; INTRAVENOUS

PALONOSETRON HYDROCHLORIDE

+ FRESENIUS KABI USA	EQ 0.25MG BASE/5ML (EQ 0.05MG BASE/ML)	N208109	001	Nov 21, 2017
----------------------	--	---------	-----	--------------

HIKMA	EQ 0.25MG BASE/2ML (EQ 0.125MG BASE/ML)	N207963	001	Aug 22, 2016
-------	---	---------	-----	--------------

POSFREA

+ AVYXA HOLDINGS	EQ 0.075MG BASE/1.5ML (EQ 0.05MG	N203050	001	Mar 01, 2016
------------------	----------------------------------	---------	-----	--------------

	BASE/ML)			
--	----------	--	--	--

PAMIDRONATE DISODIUM

INJECTABLE; INJECTION

ARELIA

+ NOVARTIS	30MG/VIAL **	N020036	001	Oct 31, 1991
------------	--------------	---------	-----	--------------

	60MG/VIAL	N020036	003	May 06, 1993
--	-----------	---------	-----	--------------

	90MG/VIAL	N020036	004	May 06, 1993
--	-----------	---------	-----	--------------

PAMIDRONATE DISODIUM

AESGEN	30MG/VIAL	A075594	001	May 06, 2002
--------	-----------	---------	-----	--------------

	90MG/VIAL	A075594	002	May 06, 2002
--	-----------	---------	-----	--------------

AM REGENT	30MG/10ML (3MG/ML)	A078942	001	Jul 25, 2008
-----------	--------------------	---------	-----	--------------

	90MG/10ML (9MG/ML)	A078942	002	Jul 25, 2008
--	--------------------	---------	-----	--------------

FRESENIUS KABI USA	30MG/VIAL	A075773	001	May 06, 2002
--------------------	-----------	---------	-----	--------------

	30MG/10ML (3MG/ML)	A076207	001	May 17, 2002
--	--------------------	---------	-----	--------------

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PAMIDRONATE DISODIUM

INJECTABLE; INJECTION

PAMIDRONATE DISODIUM

	90MG/VIAL	A075773 002	May 06, 2002
	90MG/10ML (9MG/ML)	A076207 002	May 17, 2002
MN PHARMS	30MG/VIAL	A078300 001	Mar 10, 2009
	90MG/VIAL	A078300 002	Mar 10, 2009
SUN PHARMA GLOBAL	30MG/VIAL	A077703 001	Dec 24, 2008
	90MG/VIAL	A077703 002	Dec 24, 2008
TEVA PHARMS USA	30MG/10ML (3MG/ML)	A076153 001	Mar 27, 2002
	90MG/10ML (9MG/ML)	A076153 002	Mar 27, 2002

PANCURONIUM BROMIDE

INJECTABLE; INJECTION

PANCURONIUM BROMIDE

ELKINS SINN	1MG/ML	A072058 001	Mar 23, 1988
	2MG/ML	A072059 001	Mar 23, 1988
	2MG/ML	A072060 001	Mar 23, 1988
HOSPIRA	1MG/ML	A072320 001	Jan 19, 1989
	2MG/ML	A072321 001	Jan 19, 1989
IGI LABS INC	1MG/ML	A072210 001	Mar 31, 1988
	2MG/ML	A072211 001	Mar 31, 1988
	2MG/ML	A072212 001	Mar 31, 1988
	2MG/ML	A072213 001	Mar 31, 1988
PAVULON			
+ ORGANON USA INC	1MG/ML **	N017015 002	
+	2MG/ML **	N017015 001	

PANOBINOSTAT LACTATE

CAPSULE; ORAL

FARYDAK

+ SECURA	EQ 10MG BASE	N205353 001	Feb 23, 2015
+	EQ 15MG BASE	N205353 002	Feb 23, 2015
+	EQ 20MG BASE	N205353 003	Feb 23, 2015

PANTOPRAZOLE SODIUM

INJECTABLE; INTRAVENOUS

PANTOPRAZOLE SODIUM

NORVIUM BIOSCIENCE	EQ 40MG BASE/VIAL	A208580 001	May 04, 2018
SANDOZ	EQ 40MG BASE/VIAL	A090296 001	Jul 14, 2015
SUN PHARM	EQ 40MG BASE/VIAL	A077674 001	Aug 19, 2019
TABLET, DELAYED RELEASE; ORAL			
PANTOPRAZOLE SODIUM			
JUBILANT GENERICS	EQ 20MG BASE	A090901 001	Aug 30, 2011
	EQ 40MG BASE	A090901 002	Aug 30, 2011
L PERRIGO CO	EQ 20MG BASE	A203024 001	May 07, 2014
MACLEODS PHARMS LTD	EQ 20MG BASE	A200821 001	Feb 16, 2012
	EQ 40MG BASE	A200821 002	Feb 16, 2012
MANKIND PHARMA	EQ 20MG BASE	A215880 002	Apr 25, 2024
SUN PHARM	EQ 20MG BASE	A077058 001	Sep 10, 2007
	EQ 40MG BASE	A077058 002	Sep 10, 2007
SUN PHARM INDS LTD	EQ 20MG BASE	A200794 001	May 02, 2012
	EQ 40MG BASE	A200794 002	May 02, 2012
TEVA	EQ 20MG BASE	A077056 001	Aug 02, 2007
	EQ 40MG BASE	A077056 002	Aug 02, 2007

PARAMETHADIONE

CAPSULE; ORAL

PARADIONE

ABBVIE	150MG	N006800 003	
	300MG	N006800 001	

SOLUTION; ORAL

PARADIONE

ABBVIE	300MG/ML	N006800 002	
--------	----------	-------------	--

PARAMETHASONE ACETATE

TABLET; ORAL

HALDRONE

LILLY	1MG	N012772 005	
	2MG	N012772 006	

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PARGYLINE HYDROCHLORIDE

TABLET; ORAL

EUTONYL

ABBOTT	10MG	N013448 002
	25MG	N013448 003
	50MG	N013448 004

PARICALCITOL

CAPSULE; ORAL

PARICALCITOL

LOTUS PHARM CO LTD	1MCG	A206710 001	Feb 24, 2016
	2MCG	A206710 002	Feb 24, 2016
	4MCG	A206710 003	Feb 24, 2016

ZEMPLAR

+ ABBVIE

4MCG **	N021606 003	May 26, 2005
---------	-------------	--------------

SOLUTION; INTRAVENOUS

PARICALCITOL

EPIC PHARMA LLC	0.01MG/2ML (0.005MG/ML)	A207692 001	Oct 16, 2017
HOSPIRA	0.002MG/ML (0.002MG/ML)	N201657 001	Oct 21, 2014
	0.005MG/ML (0.005MG/ML)	N201657 002	Oct 21, 2014
	0.01MG/2ML (0.005MG/ML)	N201657 003	Oct 21, 2014
RISING	0.002MG/ML (0.002MG/ML)	A203897 001	Nov 02, 2017

PAROMOMYCIN SULFATE

CAPSULE; ORAL

HUMATIN

KING PFIZER	EQ 250MG BASE	A062310 001
PARKEDALE	EQ 250MG BASE	A060521 001

PAROMOMYCIN SULFATE

SUN PHARM INDS INC	EQ 250MG BASE	A064171 001	Jun 30, 1997
--------------------	---------------	-------------	--------------

SYRUP; ORAL

HUMATIN

PARKE DAVIS	EQ 125MG BASE/5ML	A060522 001
-------------	-------------------	-------------

PAROXETINE HYDROCHLORIDE

CAPSULE; ORAL

PAXIL

+ APOTEX

EQ 10MG BASE **	N020885 001	Oct 09, 1998
-----------------	-------------	--------------

+

EQ 20MG BASE **	N020885 002	Oct 09, 1998
-----------------	-------------	--------------

+

EQ 30MG BASE **	N020885 003	Oct 09, 1998
-----------------	-------------	--------------

+

EQ 40MG BASE **	N020885 004	Oct 09, 1998
-----------------	-------------	--------------

SUSPENSION; ORAL

PAROXETINE HYDROCHLORIDE

APOTEX INC	EQ 10MG BASE/5ML	A077395 001	Dec 05, 2006
------------	------------------	-------------	--------------

PAXIL

+ APOTEX

EQ 10MG BASE/5ML	N020710 001	Jun 25, 1997
------------------	-------------	--------------

TABLET; ORAL

PAROXETINE HYDROCHLORIDE

JUBILANT GENERICS	EQ 10MG BASE	A205528 001	Nov 27, 2015
	EQ 20MG BASE	A205528 002	Nov 27, 2015
	EQ 30MG BASE	A205528 003	Nov 27, 2015
	EQ 40MG BASE	A205528 004	Nov 27, 2015
NORVIUM BIOSCIENCE	EQ 10MG BASE	A075716 001	Mar 08, 2004
	EQ 20MG BASE	A075716 002	Mar 08, 2004
	EQ 30MG BASE	A075716 003	Mar 08, 2004
	EQ 40MG BASE	A075716 004	Mar 08, 2004
ROXANE	EQ 10MG BASE	A078026 001	Jun 29, 2007
	EQ 20MG BASE	A078026 002	Jun 29, 2007
	EQ 30MG BASE	A078026 003	Jun 29, 2007
	EQ 40MG BASE	A078026 004	Jun 29, 2007
SUN PHARM INDS INC	EQ 10MG BASE	A078194 001	Jun 29, 2007
	EQ 20MG BASE	A078194 002	Jun 29, 2007
	EQ 30MG BASE	A078194 003	Jun 29, 2007
	EQ 40MG BASE	A078194 004	Jun 29, 2007
TEVA PHARMS	EQ 10MG BASE	A077082 001	Jun 29, 2007
	EQ 20MG BASE	A077082 002	Jun 29, 2007
	EQ 30MG BASE	A077082 003	Jun 29, 2007
	EQ 40MG BASE	A077082 004	Jun 29, 2007
UPSHER SMITH LABS	EQ 10MG BASE	A075566 001	Mar 08, 2004
	EQ 20MG BASE	A075566 002	Mar 08, 2004
	EQ 30MG BASE	A075566 003	Mar 08, 2004

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PAROXETINE HYDROCHLORIDE

TABLET; ORAL

PAROXETINE HYDROCHLORIDE	EQ 40MG BASE	A075566 004	Mar 08, 2004
PAXIL			
APOTEX	EQ 50MG BASE	N020031 004	Dec 29, 1992
TABLET, EXTENDED RELEASE; ORAL			
PAROXETINE HYDROCHLORIDE			
EPIC PHARMA LLC	EQ 12.5MG BASE	A213612 001	Aug 11, 2021
	EQ 25MG BASE	A213612 002	Aug 11, 2021
	EQ 37.5MG BASE	A213612 003	May 26, 2022

PAROXETINE MESYLATE

TABLET; ORAL

PEXEVA			
+ SEBELA IRELAND LTD	EQ 10MG BASE	N021299 001	Jul 03, 2003
+	EQ 20MG BASE	N021299 002	Jul 03, 2003
+	EQ 30MG BASE	N021299 003	Jul 03, 2003
+	EQ 40MG BASE	N021299 004	Jul 03, 2003

PATIROMER SORBITE X CALCIUM

POWDER; ORAL

VELTASSA			
+ VIFOR PHARMA	EQ 25.2GM BASE/PACKET	N205739 003	Oct 21, 2015

PAZOPANIB HYDROCHLORIDE

TABLET; ORAL

VOTRIENT			
+ NOVARTIS	EQ 400MG BASE **	N022465 002	Oct 19, 2009

PEGAPTANIB SODIUM

INJECTABLE; INTRAVITREAL

MACUGEN			
+ BAUSCH AND LOMB INC	EQ 0.3MG ACID/0.09ML	N021756 001	Dec 17, 2004

PEGINESATIDE ACETATE

SOLUTION; INTRAVENOUS, SUBCUTANEOUS

OMONTYS			
TAKEDA PHARMS USA	EQ 10MG BASE/ML (EQ 10MG BASE/ML)	N202799 007	Mar 27, 2012
	EQ 20MG BASE/2ML (EQ 10MG BASE/ML)	N202799 008	Mar 27, 2012
OMONTYS PRESERVATIVE FREE			
TAKEDA PHARMS USA	EQ 1MG BASE/0.5ML (EQ 1MG BASE/0.5ML)	N202799 001	Mar 27, 2012
	EQ 2MG BASE/0.5ML (EQ 2MG BASE/0.5ML)	N202799 002	Mar 27, 2012
	EQ 3MG BASE/0.5ML (EQ 3MG BASE/0.5ML)	N202799 003	Mar 27, 2012
	EQ 4MG BASE/0.5ML (EQ 4MG BASE/0.5ML)	N202799 004	Mar 27, 2012
	EQ 5MG BASE/0.5ML (EQ 5MG BASE/0.5ML)	N202799 005	Mar 27, 2012
	EQ 6MG BASE/0.5ML (EQ 6MG BASE/0.5ML)	N202799 006	Mar 27, 2012

PEMETREXED DISODIUM

POWDER; INTRAVENOUS

PEMETREXED DISODIUM			
AMNEAL	EQ 100MG BASE/VIAL	A210047 002	May 08, 2023
	EQ 500MG BASE/VIAL	A210047 001	Aug 04, 2022
BIOCON PHARMA	EQ 1GM BASE/VIAL	A211090 001	May 25, 2022
HOSPIRA	EQ 100MG BASE/VIAL	A202111 001	May 25, 2022
	EQ 500MG BASE/VIAL	A202111 002	May 25, 2022
	EQ 1GM BASE/VIAL	A202111 003	May 25, 2022
NORVIUM BIOSCIENCE	EQ 500MG BASE/VIAL	A203628 001	Mar 10, 2023

PEMETREXED DITROMETHAMINE

POWDER; INTRAVENOUS

PEMETREXED DITROMETHAMINE			
+ HOSPIRA	EQ 1GM BASE/VIAL	N208746 003	Jun 10, 2022

PEMIROLAST POTASSIUM

SOLUTION/DROPS; OPHTHALMIC

ALAMAST			
SANTEN	0.1%	N021079 001	Sep 24, 1999

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PEMOLINE

TABLET; ORAL

CYLERT

ABBOTT	18.75MG	N016832 001
	37.5MG	N016832 002
	75MG	N016832 003

PEMOLINE

ACTAVIS ELIZABETH	18.75MG	A075595 001	Feb 28, 2000
	37.5MG	A075595 002	Feb 28, 2000
	75MG	A075595 003	Feb 28, 2000
FOSUN PHARMA	18.75MG	A075286 001	Dec 27, 1999
	37.5MG	A075286 002	Jun 30, 1999
	75MG	A075286 003	Jun 30, 1999
MALLINCKRODT	18.75MG	A075726 003	Mar 30, 2001
	37.5MG	A075726 002	Mar 30, 2001
	75MG	A075726 001	Mar 30, 2001
TEVA PHARMS	18.75MG	A075030 003	Feb 22, 2000
	37.5MG	A075030 001	Jan 29, 1999
	75MG	A075030 002	Jan 29, 1999
VINTAGE PHARMS	18.75MG	A075328 001	Apr 19, 2000
	37.5MG	A075328 002	Apr 19, 2000
	75MG	A075328 003	Apr 19, 2000
WATSON LABS	18.75MG	A075287 001	Jun 13, 2001
	37.5MG	A075287 002	Sep 18, 2000
	75MG	A075287 003	Sep 18, 2000

TABLET, CHEWABLE; ORAL

CYLERT

ABBOTT	37.5MG	N017703 001
--------	--------	-------------

PEMOLINE

ACTAVIS ELIZABETH	37.5MG	A075678 001	Jul 26, 2000
TEVA PHARMS	37.5MG	A075555 001	Feb 18, 2000

PENBUTOLOL SULFATE

TABLET; ORAL

LEVATOL

+ ENDO OPERATIONS	10MG **	N018976 001	Dec 30, 1987
+	20MG **	N018976 004	Jan 05, 1989

PENICILLAMINE

CAPSULE; ORAL

CUPRIMINE

VALEANT PHARMS INTL	125MG	N019853 002
---------------------	-------	-------------

LUXZYLA

ANI PHARMS	250MG	A209921 001	May 07, 2019
------------	-------	-------------	--------------

TABLET; ORAL

PENICILLAMINE

TEVA PHARMS USA	250MG	A211497 001	Feb 13, 2020
-----------------	-------	-------------	--------------

PENICILLIN G BENZATHINE

INJECTABLE; INJECTION

BICILLIN L-A

+ KING PHARMS LLC	300,000 UNITS/ML	N050141 003
WYETH AYERST	300,000 UNITS/ML	N050131 001

PERMAPEN

CASPER PHARMA LLC	600,000 UNITS/ML	N060014 001
-------------------	------------------	-------------

SUSPENSION; ORAL

BICILLIN

WYETH AYERST	300,000 UNITS/5ML	N050126 002
--------------	-------------------	-------------

TABLET; ORAL

BICILLIN

WYETH AYERST	200,000 UNITS	N050128 001
--------------	---------------	-------------

PENICILLIN G BENZATHINE; PENICILLIN G PROCAINE

INJECTABLE; INJECTION

BICILLIN C-R

+ KING PHARMS LLC	150,000 UNITS/ML; 150,000 UNITS/ML	N050138 002
-------------------	------------------------------------	-------------

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PENICILLIN G POTASSIUM

FOR SOLUTION;ORAL

PENICILLIN

TEVA	200,000 UNITS/5ML	A060307 002
	400,000 UNITS/5ML	A060307 004

PENICILLIN G POTASSIUM

MYLAN	200,000 UNITS/5ML	A060752 003
	250,000 UNITS/5ML	A060752 002
	400,000 UNITS/5ML	A060752 001
PUREPAC PHARM	250,000 UNITS/5ML	A061740 001
	400,000 UNITS/5ML	A061740 002

PENICILLIN-2

TEVA	250,000 UNITS/5ML	A060307 003
------	-------------------	-------------

PENTIDS '200'

APOTHECON	200,000 UNITS/5ML	A062149 001
-----------	-------------------	-------------

PENTIDS '400'

APOTHECON	400,000 UNITS/5ML	A062149 002
-----------	-------------------	-------------

PFIZERPEN G

PFIZER	400,000 UNITS/5ML	A060587 001
--------	-------------------	-------------

INJECTABLE;INJECTION

PENICILLIN G POTASSIUM

APOTHECON	1,000,000 UNITS/VIAL	A060362 001
-----------	----------------------	-------------

	5,000,000 UNITS/VIAL	A060362 003
--	----------------------	-------------

	10,000,000 UNITS/VIAL	A060362 004
--	-----------------------	-------------

	20,000,000 UNITS/VIAL	A060362 002
--	-----------------------	-------------

CONSOLIDATED PHARM	500,000 UNITS/VIAL	A060806 001
--------------------	--------------------	-------------

	1,000,000 UNITS/VIAL	A060806 002
--	----------------------	-------------

	5,000,000 UNITS/VIAL	A060806 003
--	----------------------	-------------

	10,000,000 UNITS/VIAL	A060806 004
--	-----------------------	-------------

LILLY	200,000 UNITS/VIAL	A060384 004
-------	--------------------	-------------

	500,000 UNITS/VIAL	A060384 003
--	--------------------	-------------

	1,000,000 UNITS/VIAL	A060384 002
--	----------------------	-------------

	5,000,000 UNITS/VIAL	A060384 001
--	----------------------	-------------

	20,000,000 UNITS/VIAL	A060384 005
--	-----------------------	-------------

	20,000,000 UNITS/VIAL	A060601 001
--	-----------------------	-------------

PARKE DAVIS	1,000,000 UNITS/VIAL	A062003 001
-------------	----------------------	-------------

	5,000,000 UNITS/VIAL	A062003 002
--	----------------------	-------------

PFIZER	20,000,000 UNITS/VIAL	A060074 003
--------	-----------------------	-------------

SANDOZ	1,000,000 UNITS/VIAL **	A065079 001	Aug 30, 2002
--------	-------------------------	-------------	--------------

WATSON LABS INC	1,000,000 UNITS/VIAL	A062991 001	Sep 13, 1988
-----------------	----------------------	-------------	--------------

	5,000,000 UNITS/VIAL	A062991 002	Sep 13, 1988
--	----------------------	-------------	--------------

	10,000,000 UNITS/VIAL	A062991 003	Sep 13, 1988
--	-----------------------	-------------	--------------

	20,000,000 UNITS/VIAL	A062991 004	Sep 13, 1988
--	-----------------------	-------------	--------------

PFIZERPEN

PFIZER	1,000,000 UNITS/VIAL **	A060657 001
--------	-------------------------	-------------

TABLET;ORAL

PENICILLIN G POTASSIUM

APOTHECON	250,000 UNITS	A060392 003
-----------	---------------	-------------

IVAX SUB TEVA PHARMS	400,000 UNITS	A060073 004
----------------------	---------------	-------------

LILLY	250,000 UNITS	A060403 001
-------	---------------	-------------

MYLAN	200,000 UNITS	A060781 001
-------	---------------	-------------

	250,000 UNITS	A060781 002
--	---------------	-------------

	400,000 UNITS	A060781 003
--	---------------	-------------

	500,000 UNITS	A060781 005
--	---------------	-------------

	800,000 UNITS	A060781 004
--	---------------	-------------

PUREPAC PHARM	200,000 UNITS	A061588 001
---------------	---------------	-------------

	250,000 UNITS	A061588 002
--	---------------	-------------

	400,000 UNITS	A061588 003
--	---------------	-------------

TEVA	200,000 UNITS	A060306 001
------	---------------	-------------

	250,000 UNITS	A060306 002
--	---------------	-------------

	400,000 UNITS	A060306 003
--	---------------	-------------

	500,000 UNITS	A060306 004
--	---------------	-------------

WYETH AYERST	200,000 UNITS	A060413 001
--------------	---------------	-------------

	250,000 UNITS	A060413 002
--	---------------	-------------

	400,000 UNITS	A060413 003
--	---------------	-------------

PENTIDS '200'

APOTHECON	200,000 UNITS	A062155 001
-----------	---------------	-------------

PENTIDS '250'

APOTHECON	250,000 UNITS	A062155 002
-----------	---------------	-------------

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PENICILLIN G POTASSIUM

TABLET; ORAL

PENTIDS '400'

APOTHECON	400,000 UNITS	A060392 004
	400,000 UNITS	A062155 003

PENTIDS '800'

APOTHECON	800,000 UNITS	A060392 005
	800,000 UNITS	A062155 004

PFIZERPEN G

PFIZER	50,000 UNITS	A060075 001
	100,000 UNITS	A060075 002
	200,000 UNITS	A060075 003
	250,000 UNITS	A060075 004
	400,000 UNITS	A060075 005
	800,000 UNITS	A060075 006

PENICILLIN G PROCAINE

INJECTABLE; INJECTION

DURACILLIN A.S.

LILLY	300,000 UNITS/ML	A060093 001
-------	------------------	-------------

PENICILLIN G PROCAINE

CONSOLIDATED PHARM	300,000 UNITS/ML	A060800 001
	600,000 UNITS/1.2ML	A060800 002
PARKE DAVIS	300,000 UNITS/ML	A062029 001
PFIZER	300,000 UNITS/VIAL	A060099 001
	1,500,000 UNITS/VIAL	A060099 002

PFIZERPEN-AS

PFIZER	300,000 UNITS/ML	A060286 001
	600,000 UNITS/ML	A060286 002

PENICILLIN G SODIUM

INJECTABLE; INJECTION

PENICILLIN G SODIUM

BRISTOL MYERS SQUIBB	5,000,000 UNITS/VIAL	A061935 001
COPANOS	5,000,000 UNITS/VIAL	A061051 001
PHARMACIA AND UPJOHN	1,000,000 UNITS/VIAL	A061046 001

INJECTABLE; INTRAMUSCULAR, INTRAVENOUS

PENICILLIN G SODIUM

WATSON LABS INC	5,000,000 UNITS/VIAL	A063014 001	Sep 13, 1988
-----------------	----------------------	-------------	--------------

PENICILLIN V

FOR SUSPENSION; ORAL

V-CILLIN

LILLY	125MG/0.6ML	A060002 001
-------	-------------	-------------

PENICILLIN V POTASSIUM

FOR SOLUTION; ORAL

BEEPEN-VK

GLAXOSMITHKLINE	EQ 125MG BASE/5ML	A062270 001
	EQ 250MG BASE/5ML	A062270 002

BETAPEN-VK

APOTHECON	EQ 125MG BASE/5ML	A061149 001
	EQ 250MG BASE/5ML	A061149 002

LEDERCILLIN VK

LEDERLE	EQ 125MG BASE/5ML	A060136 001
	EQ 250MG BASE/5ML	A060136 002

PEN-VEE K

WYETH AYERST	EQ 125MG BASE/5ML	A060007 001
	EQ 250MG BASE/5ML	A060007 002

PENAPAR-VK

PARKE DAVIS	EQ 125MG BASE/5ML	A062002 001
	EQ 250MG BASE/5ML	A062002 002

PENICILLIN V POTASSIUM

BELCHER PHARMS	EQ 125MG BASE/5ML	A061529 001
	EQ 250MG BASE/5ML	A061529 002

CHARTWELL RX	EQ 125MG BASE/5ML	A062981 001	Feb 10, 1989
	EQ 250MG BASE/5ML	A062981 002	Feb 10, 1989

MYLAN	EQ 125MG BASE/5ML	A061624 002
	EQ 250MG BASE/5ML	A061624 001

PUREPAC PHARM	EQ 125MG BASE/5ML	A061758 001
	EQ 250MG BASE/5ML	A061758 002

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PENICILLIN V POTASSIUM

FOR SOLUTION;ORAL

PFIZERPEN VK			
PFIZER	EQ 125MG BASE/5ML		A061815 001
	EQ 250MG BASE/5ML		A061815 002
V-CILLIN K			
LILLY	EQ 125MG BASE/5ML		A060004 001
	EQ 250MG BASE/5ML		A060004 002
VEETIDS			
APOTHECON	EQ 125MG BASE/5ML		A061410 001
	EQ 250MG BASE/5ML		A061410 002
VEETIDS '125'			
APOTHECON	EQ 125MG BASE/5ML		A061206 001
	EQ 125MG BASE/5ML		A062153 001
VEETIDS '250'			
APOTHECON	EQ 250MG BASE/5ML		A061206 002
	EQ 250MG BASE/5ML		A062153 002

TABLET;ORAL

BEEPEN-VK			
GLAXOSMITHKLINE	EQ 250MG BASE		A062273 001
	EQ 500MG BASE		A062273 002
BETAPEN-VK			
BRISTOL	EQ 250MG BASE		A061150 001
	EQ 500MG BASE		A061150 002
LEDERCILLIN VK			
LEDERLE	EQ 250MG BASE		A060134 001
	EQ 500MG BASE		A060134 002
PEN-VEE K			
WYETH AYERST	EQ 125MG BASE		A060006 001
	EQ 250MG BASE		A060006 002
	EQ 500MG BASE		A060006 003
PENAPAR-VK			
PARKE DAVIS	EQ 250MG BASE		A062001 001
	EQ 500MG BASE		A062001 002
PENICILLIN V POTASSIUM			
BELCHER PHARMS	EQ 250MG BASE		A061528 001
	EQ 500MG BASE		A061528 002
IVAX SUB TEVA PHARMS	EQ 125MG BASE		A060518 001
	EQ 250MG BASE		A060518 002
	EQ 500MG BASE		A060518 003
MYLAN	EQ 250MG BASE		A061530 001
	EQ 500MG BASE		A061530 002
PUREPAC PHARM	EQ 125MG BASE		A061571 001
	EQ 250MG BASE		A061571 002
	EQ 500MG BASE		A061571 003
SANDOZ	EQ 250MG BASE		A064071 001 Nov 30, 1995
	EQ 500MG BASE		A064071 002 Nov 30, 1995
PFIZERPEN VK			
PFIZER	EQ 250MG BASE		A061836 001
	EQ 500MG BASE		A061836 002
UTICILLIN VK			
PHARMACIA AND UPJOHN	EQ 250MG BASE		A061651 001
	EQ 500MG BASE		A061651 002
V-CILLIN K			
LILLY	EQ 125MG BASE **		A060003 001
	EQ 250MG BASE **		A060003 002
	EQ 500MG BASE **		A060003 003
VEETIDS			
APOTHECON	EQ 250MG BASE		A061411 001
	EQ 500MG BASE		A061411 002
VEETIDS '250'			
APOTHECON	EQ 250MG BASE		A061164 001
	EQ 250MG BASE		A062156 002
VEETIDS '500'			
APOTHECON	EQ 500MG BASE		A061164 002
	EQ 500MG BASE		A062156 001

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PENTAGASTRIN

INJECTABLE; INJECTION

PEPTAVLON

+ WYETH AYERST 0.25MG/ML ** N017048 001

PENTAMIDINE ISETHIONATE

FOR SOLUTION; INHALATION

NEBUPENT

FRESENIUS KABI USA 600MG/VIAL N019887 002 Mar 22, 1996

INJECTABLE; INJECTION

PENTACARINAT

ARMOUR PHARM 300MG/VIAL A073447 001 Apr 28, 1994

PENTAMIDINE ISETHIONATE

BAXTER HLTHCARE 300MG/VIAL A073617 001 Dec 18, 1995

HOSPIRA 300MG/VIAL A073479 001 Jun 30, 1992

WATSON LABS 300MG/VIAL A074303 001 Aug 17, 1995

PENTAZOCINE HYDROCHLORIDE

TABLET; ORAL

TALWIN 50

SANOFI AVENTIS US EQ 50MG BASE N016732 001

PENTAZOCINE LACTATE

INJECTABLE; INJECTION

TALWIN

+ HOSPIRA EQ 30MG BASE/ML N016194 001

PENTETATE CALCIUM TRISODIUM

SOLUTION; INHALATION, INTRAVENOUS

PENTETATE CALCIUM TRISODIUM

+ HAMELN EQ 1GM BASE/5ML (EQ 200MG BASE/ML) ** N021749 001 Aug 11, 2004

PENTETATE CALCIUM TRISODIUM YB-169

INJECTABLE; INJECTION

YTTERBIUM YB 169 DTPA

3M 2mCi/ML N017518 001

PENTETATE ZINC TRISODIUM

SOLUTION; INHALATION, INTRAVENOUS

PENTETATE ZINC TRISODIUM

+ HAMELN PHARMA PLUS EQ 1GM BASE/5ML (EQ 200MG BASE/ML) ** N021751 001 Aug 11, 2004

PENTOBARBITAL

ELIXIR; ORAL

NEMBUTAL

SCIEGEN PHARMS INC 18.2MG/5ML A083244 001

PENTOBARBITAL SODIUM

CAPSULE; ORAL

NEMBUTAL SODIUM

EPIC PHARMA LLC 30MG A084095 001

50MG A084093 001

100MG A083245 001

PENTOBARBITAL SODIUM

LANNETT 50MG A085937 001

100MG A085915 001

VITARINE 100MG A083284 001

WHITEWORTH TOWN PLSN 100MG A083338 001

SODIUM PENTOBARBITAL

ANABOLIC 100MG A084590 001

ELKINS SINN 100MG A083368 001

EVERYLIFE 100MG A083259 001

HALSEY 100MG A084677 001

IVAX SUB TEVA PHARMS 50MG A083461 001

100MG A083461 002

PARKE DAVIS 100MG A084156 001

PERRIGO 100MG A084560 001

PUREPAC PHARM 100MG A083301 001

VALEANT PHARM INTL 100MG A083264 001

WATSON LABS 100MG A085791 001

WYETH AYERST 100MG A083239 001

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PENTOBARBITAL SODIUM

INJECTABLE; INJECTION

PENTOBARBITAL SODIUM

ELKINS SINN 50MG/ML A083270 001

SODIUM PENTOBARBITAL

WYETH AYERST 50MG/ML A083261 001

SUPPOSITORY; RECTAL

NEMBUTAL

SCIEGEN PHARMS INC 30MG A083247 001 Jan 25, 1982

60MG A083247 002 Jan 25, 1982

120MG A083247 003 Jan 25, 1982

200MG A083247 004 Jan 25, 1982

TABLET; ORAL

PENTOBARBITAL SODIUM

VITARINE 100MG A083285 001

SODIUM PENTOBARBITAL

NEXGEN PHARMA INC 100MG A084238 001

PENTOLINIUM TARTRATE

INJECTABLE; INJECTION

ANSOLYSEN

WYETH AYERST 10MG/ML N009372 001

PENTOSTATIN

INJECTABLE; INJECTION

PENTOSTATIN

RISING 10MG/VIAL A203554 001 Sep 19, 2014

PENTOXIFYLLINE

TABLET, EXTENDED RELEASE; ORAL

PENTOXIFYLLINE

ANI PHARMS 400MG A075107 001 Sep 04, 1998

400MG A075199 001 Sep 03, 1999

HERITAGE 400MG A074877 001 Jul 08, 1997

IMPAX LABS 400MG A075093 001 Aug 10, 1999

PLIVA 400MG A074874 001 May 25, 1999

PENTOXIL

UPSHER SMITH LABS 400MG A074962 001 Mar 31, 1999

TRENAL

+ VALIDUS PHARMS 400MG ** N018631 001 Aug 30, 1984

PERFLUBRON

LIQUID; ORAL

IMAGENT

ALLIANCE PHARM 100% N020091 001 Aug 13, 1993

PERFLUOROPOLYMETHYLISOPROPYL ETHER; POLYTETRAFLUOROETHYLENE

PASTE; TOPICAL

SKIN EXPOSURE REDUCTION PASTE AGAINST CHEMICAL WARFARE AGENTS

US ARMY MED RES 50%;50% N021084 001 Feb 17, 2000

PERGOLIDE MESYLATE

TABLET; ORAL

PERGOLIDE MESYLATE

IVAX SUB TEVA PHARMS EQ 0.05MG BASE A076094 001 Sep 04, 2003

EQ 0.25MG BASE A076094 002 Sep 04, 2003

EQ 1MG BASE A076094 003 Sep 04, 2003

STRIDES PHARMA EQ 0.05MG BASE A076061 001 Nov 27, 2002

EQ 0.25MG BASE A076061 002 Nov 27, 2002

EQ 1MG BASE A076061 003 Nov 27, 2002

PERMAX

VALEANT PHARM INTL EQ 0.05MG BASE N019385 001 Dec 30, 1988

EQ 0.25MG BASE N019385 002 Dec 30, 1988

EQ 1MG BASE N019385 003 Dec 30, 1988

PERINDOPRIL ERBUMINE

TABLET; ORAL

ACEON

+ SYMPLMED PHARMS LLC 2MG N020184 001 Dec 30, 1993

+ 4MG N020184 002 Dec 30, 1993

+ 8MG N020184 003 Dec 30, 1993

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PERINDOPRIL ERBUMINE

TABLET; ORAL

PERINDOPRIL ERBUMINE

ANI PHARMS	2MG	A078138 001	Nov 10, 2009
	4MG	A078138 002	Nov 10, 2009
	8MG	A078138 003	Nov 10, 2009
APOTEX	2MG	A090463 001	Aug 30, 2010
	4MG	A090463 002	Aug 30, 2010
	8MG	A090463 003	Aug 30, 2010
HIKMA	2MG	A090072 001	Nov 10, 2009
	4MG	A090072 002	Nov 10, 2009
	8MG	A090072 003	Nov 10, 2009
LUPIN LTD	2MG	A078263 001	Jan 27, 2010
	4MG	A078263 002	Jan 27, 2010
	8MG	A078263 003	Jan 27, 2010

PERMETHRIN

CREAM; TOPICAL

PERMETHRIN

DR REDDYS LABS EU	5%	A209732 001	Aug 01, 2023
-------------------	----	-------------	--------------

LOTION; TOPICAL

NIX

GLAXOSMITHKLINE	1% **	N019435 001	Mar 31, 1986
-----------------	-------	-------------	--------------

PERPHENAZINE

CONCENTRATE; ORAL

PERPHENAZINE

PHARM ASSOC	16MG/5ML	A040360 001	May 25, 2001
-------------	----------	-------------	--------------

TRILAFON

SCHERING	16MG/5ML	N011557 001	
----------	----------	-------------	--

INJECTABLE; INJECTION

TRILAFON

SCHERING	5MG/ML	N011213 002	
----------	--------	-------------	--

SYRUP; ORAL

TRILAFON

SCHERING	2MG/5ML	N011294 002	
----------	---------	-------------	--

TABLET; ORAL

PERPHENAZINE

ANI PHARMS	2MG **	A089707 001	Sep 10, 1987
	4MG **	A089708 001	Sep 10, 1987
	8MG	A089456 001	Sep 10, 1987
	16MG	A089457 001	Sep 10, 1987
MYLAN	2MG	A206691 001	Apr 14, 2017
	4MG	A206691 002	Apr 14, 2017
	8MG	A206691 003	Apr 14, 2017
	16MG	A206691 004	Apr 14, 2017

TRILAFON

+ SCHERING	2MG **	N010775 001	
------------	--------	-------------	--

+ SCHERING	4MG **	N010775 002	
------------	--------	-------------	--

+ SCHERING	8MG **	N010775 003	
------------	--------	-------------	--

+ SCHERING	16MG **	N010775 004	
------------	---------	-------------	--

TABLET, EXTENDED RELEASE; ORAL

TRILAFON

SCHERING	8MG	N011361 002	
----------	-----	-------------	--

PEXIDARTINIB HYDROCHLORIDE

CAPSULE; ORAL

TURALIO

+ DAIICHI SANKYO INC	EQ 200MG BASE	N211810 001	Aug 02, 2019
----------------------	---------------	-------------	--------------

PHENACEMIDE

TABLET; ORAL

PHENURONE

+ ABBVIE	500MG **	N007707 001	
----------	----------	-------------	--

PHENAZOPYRIDINE HYDROCHLORIDE; SULFAMETHOXAZOLE

TABLET; ORAL

AZO GANTANOL

+ ROCHE	100MG; 500MG **	N013294 001	Sep 10, 1987
---------	-----------------	-------------	--------------

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

<u>PHENAZOPYRIDINE HYDROCHLORIDE; SULFAMETHOXAZOLE; TRIMETHOPRIM</u>			
TABLET; ORAL			
SULFAMETHOXAZOLE AND TRIMETHOPRIM AND PHENAZOPYRIDINE HYDROCHLORIDE			
ABLE	200MG, N/A, N/A; N/A, 800MG, 160MG	N021105 001	Jun 26, 2001
<u>PHENAZOPYRIDINE HYDROCHLORIDE; SULFISOXAZOLE</u>			
TABLET; ORAL			
AZO GANTRISIN			
+ ROCHE	50MG; 500MG **	N019358 001	Aug 31, 1990
<u>PHENDIMETRAZINE TARTRATE</u>			
CAPSULE; ORAL			
PHENAZINE			
MAST MM	35MG	A086523	001
	35MG	A086524	001
	35MG	A086525	001
PHENDIMETRAZINE TARTRATE			
SANDOZ	35MG	A085633	001
	35MG	A085694	001
	35MG	A085702	001
VIRTUS	35MG	A085695	001
VITARINE	35MG	A085634	001
	35MG	A085645	001
	35MG	A085670	001
	35MG	A086403	001
	35MG	A086408	001
	35MG	A086410	001
	35MG	A087424	001
SPRX-3			
SOLVAY	35MG	A085897	001
STATOBEX			
TEVA	35MG	A085507	001
X-TROZINE			
SHIRE RICHWOOD	35MG	A087394	001 Sep 22, 1982
CAPSULE, EXTENDED RELEASE; ORAL			
BONTRIL			
VALEANT	105MG	A088021	001 Sep 21, 1982
MELFIAT-105			
NUMARK	105MG	A087487	001 Oct 13, 1982
PHENDIMETRAZINE TARTRATE			
GRAHAM DM	105MG	A087214	001 May 26, 1982
	105MG	A088020	001 Aug 16, 1982
	105MG	A088028	001 Aug 16, 1982
	105MG	A088062	001 Sep 13, 1982
	105MG	A088063	001 Sep 10, 1982
	105MG	A088111	001 Oct 18, 1982
VIRTUS	105MG	A087378	001
SPRX-105			
NUMARK	105MG	A088024	001 Dec 22, 1982
X-TROZINE L.A.			
SHIRE RICHWOOD	105MG	A087371	001 Aug 24, 1982
TABLET; ORAL			
ADPHEN			
FERNDAL LABS	35MG	A083655	001
ALPHAZINE			
SANDOZ	35MG	A085034	001
CAM-METRAZINE			
ABC HOLDING	35MG	A085511	001
CAMALL	35MG	A085756	001
CHARTWELL RX	35MG	A083922	001
	35MG	A085318	001
	35MG	A085320	001
	35MG	A085321	001
DI-METREX			
PVT FORM	35MG	A085698	001
MELFIAT			
NUMARK	35MG	A083790	002
METRA			
FOREST PHARMS	35MG	A083754	001

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PHENDIMETRAZINE TARTRATE

TABLET; ORAL

PHENAZINE

MAST MM 35MG A087305 001

PHENAZINE-35

ABC HOLDING 35MG A085512 001

PHENDIMETRAZINE TARTRATE

BARR 35MG A083644 001

35MG A083684 001

35MG A083686 001

35MG A083687 001

35MG A084831 001

35MG A084834 001

35MG A084835 001

CHARTWELL RX 35MG A085761 001

35MG A085941 001 Jun 27, 1983

FERNDALE LABS 35MG A086834 001 Sep 15, 1983

INWOOD LABS 35MG A084740 001

35MG A084741 001

35MG A084742 001

35MG A084743 001

IVAX PHARMS 35MG A085611 001

35MG A085612 001

IVAX SUB TEVA PHARMS 35MG A083682 001

KV PHARM 35MG A084138 001

35MG A084141 001

35MG A085525 001

MFG CHEMISTS 35MG A085914 001

NEXGEN PHARMA INC 35MG A086020 001

NOSTRUM LABS INC 35MG A203600 001 Dec 27, 2017

NUMARK 35MG A083790 001

PVT FORM 35MG A085199 001

35MG A085697 001

SANDOZ 35MG A085402 001

35MG A085830 001

35MG A086370 001

SOLVAY 35MG A083993 001

UPSHER SMITH LABS 35MG A084399 001

USL PHARMA 35MG A083805 001

35MG A084398 001

VIRTUS 35MG A085497 001

35MG A086365 001

VITARINE 35MG A085519 001

35MG A086005 001

35MG A086106 001

WATSON LABS 35MG A085767 001

35MG A085768 001

35MG A085770 001

35MG A085773 001

PLEGINE

WYETH AYERST 35MG ** N012248 001

STATOBEX

TEVA 35MG A086013 001

STATOBEX-G

TEVA 35MG A085095 001

X-TROZINE

SHIRE RICHWOOD 35MG A086550 001

35MG A086551 001

35MG A086552 001

35MG A086553 001

35MG A086554 001

PHENINDIONE

TABLET; ORAL

HEDULIN

SANOFI AVENTIS US 50MG N008767 002

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PHENMETRAZINE HYDROCHLORIDE

TABLET;ORAL

PRELUDIN

BOEHRINGER INGELHEIM 25MG N010460 005

TABLET, EXTENDED RELEASE;ORAL

PRELUDIN

BOEHRINGER INGELHEIM 50MG N011752 004

75MG N011752 003

PHENOXYBENZAMINE HYDROCHLORIDE

CAPSULE;ORAL

PHENOXYBENZAMINE HYDROCHLORIDE

HIKMA 10MG A201050 001 Jul 16, 2012

PHENPROCOUMON

TABLET;ORAL

LIQUAMAR

ORGANON USA INC 3MG N011228 001

PHENSUXIMIDE

CAPSULE;ORAL

MILONTIN

PARKE DAVIS 500MG N008855 004

PHENTERMINE HYDROCHLORIDE

CAPSULE;ORAL

ADIPEX-P

+ TEVA 37.5MG A088023 001 Aug 02, 1983

FASTIN

GLAXOSMITHKLINE 30MG ** N017352 001

OBESTIN-30

FERNDALE LABS 30MG A087144 001

OBY-TRIM

SHIRE RICHWOOD 30MG A087764 001 Mar 18, 1982

ONA-MAST

MAST MM 30MG A086511 001

30MG A086516 001

PHENTERMINE HYDROCHLORIDE

ABC HOLDING 30MG A085411 001

ABLE 15MG A040497 001 Mar 13, 2003

30MG A040403 001 Aug 30, 2001

30MG A040427 001 Aug 30, 2001

BARR 15MG A090591 001 Mar 18, 2010

30MG A090591 002 Mar 18, 2010

CAMALL 15MG A086735 001

30MG A087226 001

CHARTWELL RX 18.75MG A088576 001 May 23, 1984

30MG A085417 001

30MG A086732 002

30MG A087215 001

37.5MG A087915 001 Dec 22, 1983

37.5MG A087918 001 Dec 22, 1983

37.5MG A087930 001 Oct 14, 1983

37.5MG A088610 001 Jun 04, 1984

37.5MG A088611 001 Jun 04, 1984

37.5MG A088625 001 Aug 23, 1984

DURAMED PHARMS BARR 30MG A088948 001 Apr 25, 1986

ELITE LABS INC 15MG A040460 001 Jan 14, 2003

30MG A040227 001 Jun 18, 1997

30MG A040448 001 Jan 22, 2003

37.5MG A040228 001 Jun 19, 1997

INVAGEN PHARMS 15MG A202858 001 Feb 14, 2014

30MG A202858 002 Feb 14, 2014

30MG A204414 001 May 05, 2014

37.5MG A202846 001 Feb 05, 2014

IVAX PHARMS 30MG A086329 001

LANNETT 15MG A087022 002 Jan 20, 2012

30MG A087022 001 Feb 03, 1983

LANNETT CO INC 30MG A091359 001 Jul 16, 2010

37.5MG A201961 001 Jul 20, 2011

SANDOZ 30MG A087208 001

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PHENTERMINE HYDROCHLORIDE

CAPSULE;ORAL

PHENTERMINE HYDROCHLORIDE

	30MG	A087223	001	
	37.5MG	A088414	001	Oct 19, 1983
SUN PHARM INDUSTRIES	30MG	A040525	001	Oct 23, 2003
	37.5MG	A040527	001	Oct 23, 2003
TEVA	30MG	A086911	001	
	30MG	A087126	001	
	30MG	A087777	001	Nov 01, 1985
	30MG	A088612	001	Apr 04, 1984
	30MG	A088613	001	Apr 09, 1984
	30MG	A088614	001	Apr 09, 1984
TG UNITED INC	30MG	A040083	001	Mar 07, 1997
UPSHER SMITH LABS	30MG	A084487	001	Apr 09, 1982
	30MG	A088430	001	Mar 27, 1984
USL PHARMA	30MG	A088797	001	Dec 10, 1984
VITARINE	30MG	A087202	001	
	30MG	A087235	001	
WATSON LABS	30MG	A086740	001	Mar 21, 1985

TABLET;ORAL

ONA-MAST

MAST MM

8MG

A086260 001

PHENTERMINE HYDROCHLORIDE

ABLE

37.5MG

A040402 001 Aug 30, 2001

ACTAVIS ELIZABETH

37.5MG

A040276 001 Nov 25, 1998

BARR

37.5MG

A090470 001 Aug 31, 2009

CHARTWELL RX

8MG

A083923 001

8MG

A085319 001

37.5MG

A087805 001 Dec 06, 1982

37.5MG

A088596 001 Apr 04, 1984

INVAGEN PHARMS

37.5MG

A202942 001 Feb 05, 2014

IVAX PHARMS

8MG

A085553 001

LANNETT

37.5MG

A040555 001 Apr 15, 2005

NOVAST LABS

37.5MG

A091451 001 Sep 21, 2012

SANDOZ

8MG

A085671 001

8MG

A085689 001

SANDOZ INC

30MG

A088605 001 Sep 28, 1987

SUN PHARM INDS INC

37.5MG

A040790 001 Aug 21, 2007

+

USL PHARMA

8MG

A083804 001

37.5MG

A088910 001 Jul 17, 1985

37.5MG

A088917 001 Jul 17, 1985

VITARINE

8MG

A086453 001

8MG

A086456 001

WATSON LABS

8MG

A085739 001

TORA

SOLVAY

8MG

A084035 001

WILPO

+

SANDOZ

8MG **

N012737 001

TABLET, ORALLY DISINTEGRATING;ORAL

SUPRENZA

CITIUS PHARMS

15MG **

N202088 001 Jun 13, 2011

30MG **

N202088 002 Jun 13, 2011

37.5MG **

N202088 003 Mar 27, 2012

PHENTERMINE RESIN COMPLEX

CAPSULE, EXTENDED RELEASE;ORAL

IONAMIN

UCB INC

EQ 15MG BASE **

N011613 004

EQ 30MG BASE **

N011613 002

PHENTERMINE RESIN 30

QUANTUM PHARMICS

EQ 30MG BASE

A089120 001 Feb 04, 1988

PHENTERMINE RESIN COMPLEX

LANNETT CO INC

EQ 15MG BASE

A040872 001 Jul 28, 2011

EQ 30MG BASE

A040872 002 Jul 28, 2011

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PHENTOLAMINE MESYLATE

INJECTABLE; INJECTION

REGITINE

+ NOVARTIS

5MG/VIAL **

N008278 003

PHENYL AMINOSALICYLATE

POWDER; ORAL

PHENY-PAS-TEBAMIN

PHARM RES ASSOC

50%

N011695 002

TABLET; ORAL

PHENY-PAS-TEBAMIN

PHARM RES ASSOC

500MG

N011695 003

PHENYLBUTAZONE

CAPSULE; ORAL

AZOLID

SANOFI AVENTIS US

100MG

A087260 001

BUTAZOLIDIN

NOVARTIS

100MG

N008319 009

PHENYLBUTAZONE

CHARTWELL RX

100MG

A087774 001 Jun 16, 1982

IVAX PHARMS

100MG

A088218 001 Jun 24, 1983

SUN PHARM INDUSTRIES

100MG

A088994 001 Dec 04, 1985

WATSON LABS

100MG

A087756 001 Dec 17, 1982

TABLET; ORAL

AZOLID

SANOFI AVENTIS US

100MG

A087091 001

BUTAZOLIDIN

NOVARTIS

100MG

N008319 008

PHENYLBUTAZONE

FOSUN PHARMA

100MG

A084339 001

SUN PHARM INDUSTRIES

100MG

A088863 001 Dec 04, 1985

WATSON LABS

100MG

A086151 001

100MG

A087674 001 Apr 21, 1982

PHENYLEPHRINE HYDROCHLORIDE

SOLUTION; INTRAVENOUS

BIORPHEN

+ DR REDDYS LABS SA

10MG/ML (10MG/ML)

N212909 002 Mar 11, 2021

PHENYLEPHRINE HYDROCHLORIDE

ACCORD HLTHCARE

10MG/ML (10MG/ML)

A213237 001 Jul 01, 2020

50MG/5ML (10MG/ML)

A213237 002 Jul 01, 2020

100MG/10ML (10MG/ML)

A213237 003 Jul 01, 2020

ENDO OPERATIONS

10MG/ML (10MG/ML)

A210025 001 Dec 21, 2018

HAINAN POLY

10MG/ML (10MG/ML)

A218412 001 Mar 14, 2024

50MG/5ML (10MG/ML)

A218412 002 Mar 14, 2024

100MG/10ML (10MG/ML)

A218412 003 Mar 14, 2024

PHENYLEPHRINE HYDROCHLORIDE; PROMETHAZINE HYDROCHLORIDE

SYRUP; ORAL

PHENERGAN VC

+ ANI PHARMS

5MG/5ML; 6.25MG/5ML **

N008604 003 Apr 02, 1984

PHENYLEPHRINE HYDROCHLORIDE AND PROMETHAZINE HYDROCHLORIDE

HIKMA

5MG/5ML; 6.25MG/5ML

A040675 001 Dec 23, 2014

PHERAZINE VC

HALSEY

5MG/5ML; 6.25MG/5ML

A088868 001 Mar 02, 1987

PROMETH VC PLAIN

+ G AND W LABS INC

5MG/5ML; 6.25MG/5ML

A088761 001 Nov 08, 1984

PROMETHAZINE VC PLAIN

CENCI

5MG/5ML; 6.25MG/5ML

A088815 001 Nov 22, 1985

XTTRIUM LABS INC

5MG/5ML; 6.25MG/5ML

A088897 001 Jan 04, 1985

PHENYLEPHRINE HYDROCHLORIDE; PYRILAMINE MALEATE

SOLUTION/DROPS; OPHTHALMIC

PREFRIN-A

ALLERGAN

0.12%; 0.1%

N007953 001

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PHENYLEPHRINE HYDROCHLORIDE; TROPICAMIDE

SPRAY, METERED;OPHTHALMIC

MYDCOMBI

+ EYENOVIA 2.5%;1% N215352 001 May 05, 2023

PHENYTOIN

SUSPENSION;ORAL

DILANTIN-30

VIATRIS 30MG/5ML N008762 002

PHENYTOIN

ACTAVIS MID ATLANTIC 125MG/5ML A089892 001 Sep 25, 1992

PAI HOLDINGS PHARM 125MG/5ML A040420 001 Apr 19, 2002

VISTAPHARM 125MG/5ML A040342 001 Jan 31, 2001

125MG/5ML A040342 002 Aug 18, 2005

PHENYTOIN SODIUM

CAPSULE;ORAL

DIPHENYLAN SODIUM

CHARTWELL MOLECULAR 30MG PROMPT A080857 001

100MG PROMPT A080857 002

EXTENDED PHENYTOIN SODIUM

ANI PHARMS 100MG EXTENDED A040435 001 Jun 20, 2003

100MG EXTENDED A089441 001 Dec 18, 1986

LUPIN LTD 100MG EXTENDED A211633 001 Sep 30, 2019

MYLAN 100MG EXTENDED A040298 001 Dec 28, 1998

SUN PHARM INDS 200MG EXTENDED A040731 001 Jun 30, 2008

300MG EXTENDED A040731 002 Jun 30, 2008

SUN PHARM INDS (IN) 100MG EXTENDED A040621 001 Dec 11, 2006

UNICHEM 100MG EXTENDED A213834 001 Oct 13, 2022

WOCKHARDT 30MG EXTENDED A040759 001 Dec 18, 2007

WOCKHARDT USA 100MG EXTENDED A040732 001 Jan 30, 2008

PHENYTEX

WATSON LABS 100MG EXTENDED A088711 001 Dec 21, 1984

PHENYTOIN SODIUM

PHARMERAL 100MG PROMPT A085435 001

WATSON LABS 100MG PROMPT A085894 001

PROMPT PHENYTOIN SODIUM

ANI PHARMS 100MG PROMPT A080259 001

WATSON LABS 100MG PROMPT A080905 001

INJECTABLE;INJECTION

DILANTIN

PARKE DAVIS 50MG/ML N010151 001

PHENYTOIN SODIUM

AM REGENT 50MG/ML A040781 001 Dec 04, 2007

FRESENIUS KABI USA 50MG/ML A089003 001 May 31, 1985

HOSPIRA 50MG/ML A089521 001 Mar 17, 1987

50MG/ML A089744 001 Dec 18, 1987

MARSAM PHARMS LLC 50MG/ML A089501 001 Oct 13, 1987

50MG/ML A089779 001 Nov 27, 1992

SMITH AND NEPHEW 50MG/ML A088519 001 Dec 19, 1984

50MG/ML A088521 001 Dec 18, 1984

SOLOPAK 50MG/ML A088520 001 Dec 17, 1984

WARNER CHILCOTT 50MG/ML A089900 001 Mar 30, 1990

WATSON LABS 50MG/ML A085434 001

PHYTONADIONE

INJECTABLE;INJECTION

AQUAMEPHYTON

+ PAI HOLDINGS PHARM 1MG/0.5ML ** N012223 002

+ 10MG/ML ** N012223 001

KONAKION

ROCHE 1MG/0.5ML N011745 001

10MG/ML N011745 003

PHYTONADIONE

CIPLA 1MG/0.5ML A212424 001 Apr 22, 2022

10MG/ML A212424 002 Apr 22, 2022

GLAXOSMITHKLINE 1MG/0.5ML A084060 001

10MG/ML A084060 002

VITAMIN K1

HOSPIRA 10MG/ML A087956 001 Jul 25, 1983

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PHYTONADIONE

TABLET;ORAL

MEPHYTON

+ BAUSCH 5MG ** N010104 003

PILOCARPINE

INSERT, EXTENDED RELEASE;OPHTHALMIC

OCUSERT PILO-20

EPIC PHARMA LLC 5MG N017431 001

OCUSERT PILO-40

EPIC PHARMA LLC 11MG N017548 001

PILOCARPINE HYDROCHLORIDE

GEL;OPHTHALMIC

PILOPINE HS

ALCON 4% N018796 001 Oct 01, 1984

PIMAVANSERIN TARTRATE

CAPSULE;ORAL

PIMAVANSERIN

MSN EQ 34MG BASE A214925 001 Jan 16, 2024

ZYDUS EQ 34MG BASE A214493 001 Jan 16, 2024

TABLET;ORAL

NUPLAZID

+ ACADIA PHARMS INC EQ 17MG BASE ** N207318 001 Apr 29, 2016

PIMAVANSERIN

ZYDUS EQ 10MG BASE A214502 001 Jan 16, 2024

PIMOZIDE

TABLET;ORAL

ORAP

+ TEVA 1MG ** N017473 003 Aug 27, 1997

+ 2MG ** N017473 001 Jul 31, 1984

PINACIDIL

CAPSULE, EXTENDED RELEASE;ORAL

PINDAC

LEO PHARM 12.5MG N019456 001 Dec 28, 1989

25MG N019456 002 Dec 28, 1989

PINDOLOL

TABLET;ORAL

PINDOLOL

COSETTE 5MG A073661 001 Oct 31, 1993

5MG A073687 001 Feb 26, 1993

5MG A074123 001 Apr 17, 1997

10MG A073661 002 Oct 31, 1993

10MG A073687 002 Feb 26, 1993

10MG A074123 002 Apr 17, 1997

MYLAN PHARMS INC 5MG A074013 001 Sep 24, 1992

10MG A074018 001 Sep 24, 1992

NOSTRUM LABS 5MG A074474 001 Oct 28, 1996

10MG A074474 002 Oct 28, 1996

PUREPAC PHARM 5MG A074125 001 Apr 28, 1993

10MG A074125 002 Apr 28, 1993

WATSON LABS 5MG A074437 001 Feb 27, 1995

10MG A074437 002 Feb 27, 1995

ZYDUS PHARMS 5MG A209866 001 Aug 18, 2017

10MG A209866 002 Aug 18, 2017

VISKEN

+ NOVARTIS 5MG ** N018285 001 Sep 03, 1982

+ 10MG ** N018285 002 Sep 03, 1982

PIOGLITAZONE HYDROCHLORIDE

TABLET;ORAL

PIOGLITAZONE HYDROCHLORIDE

CHARTWELL RX EQ 15MG BASE A078383 001 Mar 12, 2013

EQ 30MG BASE A078383 002 Mar 12, 2013

EQ 45MG BASE A078383 003 Mar 12, 2013

NORVIUM BIOSCIENCE EQ 15MG BASE A076801 001 Aug 17, 2012

EQ 30MG BASE A076801 002 Aug 17, 2012

EQ 45MG BASE A076801 003 Aug 17, 2012

NOSTRUM LABS INC EQ 15MG BASE A078472 001 Feb 13, 2013

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PIOGLITAZONE HYDROCHLORIDE

TABLET; ORAL

PIOGLITAZONE HYDROCHLORIDE

	EQ 30MG BASE	A078472 002	Feb 13, 2013
	EQ 45MG BASE	A078472 003	Feb 13, 2013
TORRENT PHARMS LTD	EQ 15MG BASE	A091298 001	Feb 13, 2013
	EQ 30MG BASE	A091298 002	Feb 13, 2013
	EQ 45MG BASE	A091298 003	Feb 13, 2013

PIPECURONIUM BROMIDE

INJECTABLE; INJECTION

ARDUAN

ORGANON USA INC	10MG/VIAL	N019638 001	Jun 26, 1990
-----------------	-----------	-------------	--------------

PIPERACETAZINE

TABLET; ORAL

QUIDE

DOW PHARM	10MG	N013615 001	
	25MG	N013615 002	

PIPERACILLIN SODIUM

INJECTABLE; INJECTION

PIPRACIL

WYETH PHARMS INC	EQ 2GM BASE/VIAL	A062750 001	Oct 13, 1987
+	EQ 2GM BASE/VIAL **	N050545 002	
	EQ 3GM BASE/VIAL	A062750 002	Oct 13, 1987
+	EQ 3GM BASE/VIAL **	N050545 003	
	EQ 4GM BASE/VIAL	A062750 003	Oct 13, 1987
+	EQ 4GM BASE/VIAL **	N050545 004	
+	EQ 40GM BASE/VIAL **	N050545 006	Sep 30, 1985

PIPERACILLIN SODIUM; TAZOBACTAM SODIUM

INJECTABLE; INJECTION

PIPERACILLIN AND TAZOBACTAM

ASTRAL	EQ 2GM BASE/VIAL;EQ 250MG BASE/VIAL	A212287 001	Jul 29, 2019
	EQ 3GM BASE/VIAL;EQ 375MG BASE/VIAL	A212287 002	Jul 29, 2019
	EQ 4GM BASE/VIAL;EQ 500MG BASE/VIAL	A212287 003	Jul 29, 2019
EUGIA PHARMA	EQ 12GM BASE/VIAL;EQ 1.5GM BASE/VIAL	A217409 002	Oct 12, 2023
	EQ 36GM BASE/VIAL;EQ 4.5GM BASE/VIAL	A217409 001	Oct 12, 2023
FRESENIUS KABI USA	EQ 12GM BASE/VIAL;EQ 1.5GM BASE/VIAL	A206204 001	May 11, 2018
HOSPIRA INC	EQ 2GM BASE/VIAL;EQ 250MG BASE/VIAL	A065386 001	Sep 15, 2009
	EQ 3GM BASE/VIAL;EQ 375MG BASE/VIAL	A065386 002	Sep 15, 2009
	EQ 4GM BASE/VIAL;EQ 500MG BASE/VIAL	A065386 003	Sep 15, 2009
	EQ 36GM BASE/VIAL;EQ 4.5GM BASE/VIAL	A065446 001	Sep 15, 2009
ZOSYN			
+	WYETH PHARMS	EQ 2GM BASE/VIAL;EQ 250MG BASE/VIAL **	N050684 001
+		EQ 3GM BASE/VIAL;EQ 375MG BASE/VIAL **	N050684 002
+		EQ 4GM BASE/VIAL;EQ 500MG BASE/VIAL **	N050684 003
+		EQ 36GM BASE/VIAL;EQ 4.5GM BASE/VIAL **	N050684 004

PIPERAZINE CITRATE

SYRUP; ORAL

ANTEPAR

GLAXOSMITHKLINE	EQ 500MG BASE/5ML	N009102 001	
BRYREL			
SANOFI AVENTIS US	EQ 500MG BASE/5ML	N017796 001	
MULTIFUGE			
BLULINE	EQ 500MG BASE/5ML	N009452 001	
PIPERAZINE CITRATE			
ALPHARMA US PHARMS	EQ 500MG BASE/5ML	A080774 001	
LANNETT	EQ 500MG BASE/5ML	A080963 001	
LUITPOLD	EQ 500MG BASE/5ML	A080671 001	
VERMIDOL			
SOLVAY	EQ 500MG BASE/5ML	A080992 001	
TABLET; ORAL			
ANTEPAR			
GLAXOSMITHKLINE	EQ 500MG BASE	N009102 003	
PIPERAZINE CITRATE			
IMPAX LABS	EQ 250MG BASE	A080874 001	

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PIPERONYL BUTOXIDE; PYRETHRINS

AEROSOL; TOPICAL

RID MOUSSE

BAYER HEALTHCARE LLC 4%;EQ 0.33% BASE

N021043 001 Mar 07, 2000

PIPOBROMAN

TABLET; ORAL

VERCYTE

ABBOTT

10MG

N016245 001

25MG

N016245 002

PIRBUTEROL ACETATE

AEROSOL, METERED; INHALATION

MAXAIR

BAUSCH

EQ 0.2MG BASE/INH

N019009 001 Dec 30, 1986

EQ 0.2MG BASE/INH

N020014 001 Nov 30, 1992

PIRFENIDONE

CAPSULE; ORAL

PIRFENIDONE

APOTEX

267MG

A212687 001 Jun 09, 2023

CHARTWELL RX

267MG

A212404 001 Aug 28, 2023

TABLET; ORAL

ESBRIET

+ GENENTECH INC

534MG **

N208780 002 Jan 11, 2017

PIRFENIDONE

CHARTWELL RX

267MG

A212403 001 Aug 23, 2023

801MG

A212403 002 Aug 23, 2023

PIROXICAM

CAPSULE; ORAL

PIROXICAM

BRECKENRIDGE

10MG

A208991 001 Feb 21, 2018

20MG

A208991 002 Feb 21, 2018

CYCLE

10MG

A073651 001 Feb 26, 1993

20MG

A073651 002 Feb 26, 1993

EGIS

10MG

A074808 001 Jul 08, 1997

20MG

A074808 002 Jul 08, 1997

FLAMINGO PHARMS

10MG

A207938 001 Sep 09, 2016

20MG

A207938 002 Sep 09, 2016

HIKMA

10MG

A209256 001 Aug 11, 2017

20MG

A209256 002 Aug 11, 2017

IVAX SUB TEVA PHARMS

10MG

A074148 001 Jun 03, 1996

20MG

A074148 002 Jun 03, 1996

MYLAN

10MG

A074043 001 Sep 22, 1992

20MG

A074043 002 Sep 22, 1992

NORVIUM BIOSCIENCE

10MG

A074102 001 Jul 31, 1992

20MG

A074102 002 Jul 31, 1992

SCS

10MG

A074036 001 May 29, 1992

20MG

A074036 002 May 29, 1992

SUN PHARM INDUSTRIES

10MG

A073536 002 Jan 23, 2008

20MG

A073536 001 Mar 12, 1993

TEVA

10MG

A073637 001 Jan 28, 1994

20MG

A073638 001 Jan 28, 1994

TEVA PHARMS

10MG

A074103 001 Aug 28, 1992

20MG

A074103 002 Aug 28, 1992

WATSON LABS

10MG

A074287 001 May 16, 1996

10MG

A074460 001 Sep 29, 1995

20MG

A074287 002 May 16, 1996

20MG

A074460 002 Sep 29, 1995

PITAVASTATIN MAGNESIUM

TABLET; ORAL

ZYPITAMAG

+ MEDICURE

EQ 1MG BASE

N208379 001 Jul 14, 2017

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PITAVASTATIN SODIUM

TABLET;ORAL

NIKITA

+	LUPIN LTD	EQ 1MG BASE	N209875 001	Aug 04, 2017
+		EQ 2MG BASE	N209875 002	Aug 04, 2017
+		EQ 4MG BASE	N209875 003	Aug 04, 2017

PLERIXAFOR

SOLUTION;SUBCUTANEOUS

PLERIXAFOR

	ZYDUS PHARMS	24MG/1.2ML (20MG/ML)	A208980 001	Jul 26, 2023
--	--------------	----------------------	-------------	--------------

PLICAMYCIN

INJECTABLE;INJECTION

MITHRACIN

	PFIZER	2.5MG/VIAL	N050109 001	
--	--------	------------	-------------	--

PODOFILOX

SOLUTION;TOPICAL

PODOFILOX

	BAUSCH AND LOMB INC	0.5%	A090184 001	Jul 21, 2010
--	---------------------	------	-------------	--------------

POLIDOCANOL

SOLUTION;INTRAVENOUS

VARITHENA

+	PROVENSIS	77.5MG/7.75ML (10MG/ML)	N205098 002	Dec 21, 2017
---	-----------	-------------------------	-------------	--------------

POLYESTRADIOL PHOSPHATE

INJECTABLE;INJECTION

ESTRADURIN

	WYETH AYERST	40MG/AMP	N010753 001	
--	--------------	----------	-------------	--

POLYETHYLENE GLYCOL 3350

FOR SOLUTION;ORAL

GLYCOLAX

	LANNETT CO INC	17GM/SCOOPFUL	A076652 001	Jul 02, 2004
		17GM/PACKET	A090600 001	Oct 06, 2009
		17GM/SCOOPFUL	A090600 002	Oct 06, 2009

POLYETHYLENE GLYCOL 3350

	BRECKENRIDGE PHARM	17GM/SCOOPFUL	A077736 001	May 26, 2006
	NEXGEN PHARMA INC	17GM/SCOOPFUL	A077706 001	Sep 27, 2006
	PADDOCK LLC	17GM/SCOOPFUL	A077893 001	May 26, 2006
		17GM/SCOOPFUL	A090567 001	Oct 15, 2009
	TEVA PHARMS	17GM/SCOOPFUL	A077445 001	May 04, 2006

POLYETHYLENE GLYCOL 3350; POTASSIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE

FOR SOLUTION;ORAL

LAX-LYTE WITH FLAVOR PACKS

	L PERRIGO CO	420GM/BOT;1.48GM/BOT;5.72GM/BOT;11.2GM/BOT	A079232 001	Feb 25, 2010
--	--------------	--	-------------	--------------

PEG-3350, POTASSIUM CHLORIDE, SODIUM BICARBONATE, SODIUM CHLORIDE

	EXTROVIS	420GM/BOT;1.48GM/BOT;5.72GM/BOT;11.2GM/BOT	A090409 001	Apr 02, 2010
--	----------	--	-------------	--------------

	NOSTRUM LABS INC	420GM/BOT;1.48GM/BOT;5.72GM/BOT;11.2GM/BOT	A202060 001	Mar 08, 2017
--	------------------	--	-------------	--------------

TRILYTE

	AUROBINDO PHARMA USA	420GM/BOT;1.48GM/BOT;5.72GM/BOT;11.2GM/BOT	A076491 001	Feb 05, 2004
--	----------------------	--	-------------	--------------

POLYETHYLENE GLYCOL 3350; POTASSIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE; SODIUM SULFATE

FOR SOLUTION;ORAL

CLENZ-LYTE

	PADDOCK LLC	236GM/BOT;2.97GM/BOT;6.74GM/BOT;5.86GM/BOT;22.74GM/BOT	A090769 001	Jun 07, 2010
--	-------------	--	-------------	--------------

SOLUTION;ORAL

OCL

	HOSPIRA	6GM/100ML;75MG/100ML;168MG/100ML;146MG/100ML;1.29GM/100ML	N019284 001	Apr 30, 1986
--	---------	---	-------------	--------------

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

POLYETHYLENE GLYCOL 3350; POTASSIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE; SODIUM SULFATE ANHYDROUS

FOR SOLUTION;ORAL

COLYTE

NORVIUM BIOSCIENCE	120GM/PACKET;1.49GM/PACKET;3.36GM/PACKET;2.92GM/PACKET;11.36GM/PACKET **	N018983 005	Oct 26, 1984
	227.1GM/PACKET;2.82GM/PACKET;6.36GM/PACKET;5.53GM/PACKET;21.5GM/PACKET **	N018983 004	Oct 26, 1984
	227.1GM/BOT;2.82GM/BOT;6.36GM/BOT;5.53GM/BOT;21.5GM/BOT **	N018983 010	Jan 31, 1989
	240GM/BOT;2.98GM/BOT;6.72GM/BOT;5.84GM/BOT;22.72GM/BOT **	N018983 007	Jun 12, 1987
	360GM/PACKET;4.47GM/PACKET;10.08GM/PACKET;8.76GM/PACKET;34.08GM/PACKET **	N018983 006	Oct 26, 1984

COLYTE WITH FLAVOR PACKS

NORVIUM BIOSCIENCE	240GM/BOT;2.98GM/BOT;6.72GM/BOT;5.84GM/BOT;22.72GM/BOT	N018983 012	Oct 08, 1998
--------------------	--	-------------	--------------

COLYTE-FLAVORED

NORVIUM BIOSCIENCE	227.1GM/BOT;2.82GM/BOT;6.36GM/BOT;5.53GM/BOT;21.5GM/BOT **	N018983 008	Nov 14, 1991
	240GM/BOT;2.98GM/BOT;6.72GM/BOT;5.84GM/BOT;22.72GM/BOT **	N018983 009	Nov 14, 1991

GOLYTELY

+ BRAINTREE

	227.1GM/PACKET;2.82GM/PACKET;6.36GM/PACKET;5.53GM/PACKET;21.5GM/PACKET	N019011 002	Jun 02, 1992
--	--	-------------	--------------

PEG 3350 AND ELECTROLYTES

EXTROVIS

	236GM/BOT;2.97GM/BOT;6.74GM/BOT;5.86GM/BOT;22.74GM/BOT	A090928 001	Jan 28, 2010
--	--	-------------	--------------

POLYETHYLENE GLYCOL 3350 AND ELECTROLYTES

PADDOCK LLC

	240GM/BOT;2.98GM/BOT;6.72GM/BOT;5.84GM/BOT;22.72GM/BOT	A090712 001	Feb 25, 2010
--	--	-------------	--------------

FOR SUSPENSION;ORAL

CO-LAV

VINTAGE PHARMS	240GM/BOT;2.98GM/BOT;6.72GM/BOT;5.84GM/BOT;22.72GM/BOT	A073428 001	Jan 28, 1992
----------------	--	-------------	--------------

COLOVAGE

DYNAPHARM

	227.1GM/PACKET;2.82GM/PACKET;6.36GM/PACKET;5.53GM/PACKET;21.5GM/PACKET	A071320 001	Apr 20, 1988
--	--	-------------	--------------

E-Z-EM PREP LYTE

E Z EM

	236GM/BOT;2.97GM/BOT;6.74GM/BOT;5.86GM/BOT;22.74GM/BOT	A071278 001	Nov 21, 1988
--	--	-------------	--------------

GLYCOPREP

GOLDLINE

	236GM/BOT;2.97GM/BOT;6.74GM/BOT;5.86GM/BOT;22.74GM/BOT	A072319 001	Dec 23, 1988
--	--	-------------	--------------

GO-EVAC

VINTAGE PHARMS

	236GM/BOT;2.97GM/BOT;6.74GM/BOT;5.86GM/BOT;22.74GM/BOT	A073433 001	Apr 28, 1992
--	--	-------------	--------------

PEG-LYTE

SANDOZ

	236GM/BOT;2.97GM/BOT;6.74GM/BOT;5.86GM/BOT;22.74GM/BOT	A073098 001	Aug 31, 1993
--	--	-------------	--------------

POLYMYXIN B SULFATE

INJECTABLE;INJECTION

AEROSPORIN

GLAXOSMITHKLINE

	EQ 500,000 UNITS BASE/VIAL	A062036 001	
--	----------------------------	-------------	--

POLYMYXIN B SULFATE

HIKMA

	EQ 500,000 UNITS BASE/VIAL	A060716 001	
--	----------------------------	-------------	--

RISING

	EQ 500,000 UNITS BASE/VIAL	A090110 001	Jun 29, 2011
--	----------------------------	-------------	--------------

POWDER;FOR RX COMPOUNDING

POLY-RX

X GEN PHARMS

	100,000,000 UNITS/BOT	A061578 001	
--	-----------------------	-------------	--

POLYMYXIN B SULFATE

PADDOCK LLC

	100,000,000 UNITS/BOT	A062455 001	Jul 27, 1983
--	-----------------------	-------------	--------------

POLYMYXIN B SULFATE; TRIMETHOPRIM SULFATE

SOLUTION/DROPS;OPHTHALMIC

POLYTRIM

+ ALLERGAN

	10,000 UNITS/ML;EQ 1MG BASE/ML **	N050567 001	Oct 20, 1988
--	-----------------------------------	-------------	--------------

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

POLYTHIAZIDE

TABLET; ORAL

RENESE

PFIZER	1MG	N012845 001
	2MG	N012845 002
	4MG	N012845 003

POLYTHIAZIDE; PRAZOSIN HYDROCHLORIDE

CAPSULE; ORAL

MINIZIDE

PFIZER	0.5MG;EQ 1MG BASE	N017986 001
	0.5MG;EQ 2MG BASE	N017986 002
	0.5MG;EQ 5MG BASE	N017986 003

POLYTHIAZIDE; RESERPINE

TABLET; ORAL

RENESE-R

PFIZER	2MG;0.25MG	N013636 001
--------	------------	-------------

POMALIDOMIDE

CAPSULE; ORAL

POMALIDOMIDE

BRECKENRIDGE	1MG	A210111 001	Oct 30, 2020
	2MG	A210111 002	Oct 30, 2020
	3MG	A210111 003	Oct 30, 2020
	4MG	A210111 004	Oct 30, 2020
EUGIA PHARMA	1MG	A210249 001	Oct 30, 2020
	2MG	A210249 002	Oct 30, 2020
	3MG	A210249 003	Oct 30, 2020
	4MG	A210249 004	Oct 30, 2020
MYLAN	1MG	A210275 001	Jan 26, 2022
	2MG	A210275 002	Jan 26, 2022
	3MG	A210275 003	Jan 26, 2022
	4MG	A210275 004	Jan 26, 2022
TEVA PHARMS USA	1MG	A209956 001	May 04, 2022
	2MG	A209956 002	May 04, 2022
	3MG	A209956 003	May 04, 2022
	4MG	A209956 004	May 04, 2022

PONATINIB HYDROCHLORIDE

TABLET; ORAL

PONATINIB HYDROCHLORIDE

APOTEX	EQ 15MG BASE	A215893 001	Jul 14, 2023
	EQ 45MG BASE	A215893 002	Jul 14, 2023

POSACONAZOLE

TABLET, DELAYED RELEASE; ORAL

NOXAFIL

+ MERCK SHARP DOHME	100MG **	N205053 001	Nov 25, 2013
---------------------	----------	-------------	--------------

POSACONAZOLE

ACTAVIS LABS FL INC	100MG	A207355 001	Nov 30, 2022
---------------------	-------	-------------	--------------

POTASSIUM AMINOSALICYLATE

CAPSULE; ORAL

PASKALIUM

GLENWOOD	500MG	N009395 004
----------	-------	-------------

POWDER; ORAL

POTASSIUM AMINOSALICYLATE

HEXCEL	100%	A080098 001
--------	------	-------------

TABLET; ORAL

PASKALIUM

GLENWOOD	1GM	N009395 003
----------	-----	-------------

POTASSIUM CHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

K-LEASE

SAVAGE LABS	8MEQ	A073398 001	Jan 28, 1992
	10MEQ	A072427 001	Mar 28, 1990

MICRO-K

+ NESHER PHARMS	8MEQ **	N018238 001
-----------------	---------	-------------

MICRO-K 10

+ NESHER PHARMS	10MEQ **	N018238 002	May 14, 1984
-----------------	----------	-------------	--------------

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

POTASSIUM CHLORIDE

CAPSULE, EXTENDED RELEASE;ORAL

POTASSIUM CHLORIDE

CHARTWELL MOLECULAR	8MEQ	A204210 001	Mar 28, 2016
	10MEQ	A204210 002	Mar 28, 2016
ENDO OPERATIONS	8MEQ	A202886 001	Dec 26, 2013
	10MEQ	A202886 002	Dec 26, 2013
GLENMARK PHARMS LTD	10MEQ	A202868 001	Jan 19, 2016
NESHER PHARMS	10MEQ	A070980 001	Feb 17, 1987
TEVA	8MEQ	A073531 001	Apr 26, 1996
	10MEQ	A073532 001	Apr 26, 1996
TRIS PHARMA INC	8MEQ	A201944 001	Mar 04, 2016
	10MEQ	A201944 002	Mar 04, 2016

FOR SUSPENSION, EXTENDED RELEASE;ORAL

MICRO-K LS

+ KV PHARM	20MEQ/PACKET **	N019561 003	Aug 26, 1988
------------	-----------------	-------------	--------------

INJECTABLE;INJECTION

POTASSIUM CHLORIDE

+ ABRAXIS PHARM	2MEQ/ML	A080204 001	
	2MEQ/ML	A084290 001	
	2MEQ/ML	A086713 001	
	2MEQ/ML	A086714 001	
	2MEQ/ML	A087787 001	Apr 20, 1982
	2MEQ/ML	A087885 001	Feb 03, 1983
BAXTER HLTHCARE	2MEQ/ML	A080203 001	
	2MEQ/ML	A085499 001	
EPIC PHARMA LLC	2MEQ/ML	A088286 001	Sep 05, 1985
+ FRESENIUS KABI USA	2MEQ/ML	A080225 001	
	2MEQ/ML	A087817 001	Oct 20, 1982
	3MEQ/ML	A080225 003	
GD SEARLE LLC	1MEQ/ML	A086219 001	
	2MEQ/ML	A086219 002	
	2MEQ/ML	A086220 002	
	3MEQ/ML	A086219 003	
	3MEQ/ML	A086220 001	
	4MEQ/ML	A086219 004	
HOSPIRA	1MEQ/ML	A080205 003	
	1MEQ/ML	A083345 003	
	1.5MEQ/ML	A083345 001	
	2MEQ/ML	A083345 002	
	2.4MEQ/ML	A080205 004	
	3.2MEQ/ML	A080205 005	
INTL MEDICATION	2MEQ/ML	A083163 001	
LILLY	2MEQ/ML	N007865 002	
LUITPOLD	2MEQ/ML	A080221 001	
	2MEQ/ML	A080736 001	
	2MEQ/ML	A087584 001	
	2MEQ/ML	A087585 001	
MILES	1MEQ/ML	A080195 002	
	2MEQ/ML	A080195 001	
	3MEQ/ML	A080195 003	
	4MEQ/ML	A080195 004	
PHARMA SERVE NY	2MEQ/ML	A086297 001	
	2MEQ/ML	A087362 001	Mar 08, 1983
WATSON LABS	2MEQ/ML	A086208 001	
	2MEQ/ML	A089163 001	Mar 10, 1988
	2MEQ/ML	A089421 001	Jan 02, 1987
	3MEQ/ML	A086210 001	

POTASSIUM CHLORIDE 30MEQ IN PLASTIC CONTAINER

+ ICU MEDICAL INC	2.24GM/100ML	N020161 003	Aug 11, 1998
-------------------	--------------	-------------	--------------

SOLUTION;ORAL

POTASSIUM CHLORIDE

STRIDES PHARMA	40MEQ/15ML	A211665 001	Nov 17, 2022
TRIS PHARMA INC	20MEQ/15ML	A214076 001	Jan 26, 2022
	40MEQ/15ML	A214076 002	Jan 26, 2022

TABLET, EXTENDED RELEASE;ORAL

K+10

FUTURE PAK	10MEQ	A070999 001	Oct 22, 1987
------------	-------	-------------	--------------

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

POTASSIUM CHLORIDE

TABLET, EXTENDED RELEASE;ORAL

K+8

FUTURE PAK	8MEQ	A070998 001	Jan 25, 1993
K-TAB			
+ ABBVIE	8MEQ **	N018279 002	Aug 01, 1988
+	10MEQ **	N018279 001	
+	20MEQ **	N018279 003	Nov 25, 2013
KAON CL			
SAVAGE LABS	6.7MEQ	N017046 001	
KAON CL-10			
SAVAGE LABS	10MEQ	N017046 002	
KLOTRIX			
APOTHECON	10MEQ	N017850 001	
POTASSIUM CHLORIDE			
AMNEAL	15MEQ	A212861 002	May 08, 2020
AUROBINDO PHARMA LTD	10MEQ	A214728 001	Mar 31, 2021
	15MEQ	A214728 002	Mar 31, 2021
	20MEQ	A214728 003	Mar 31, 2021
BRECKENRIDGE	10MEQ	A213588 001	Aug 21, 2020
	20MEQ	A213588 002	Aug 21, 2020
CHARTWELL RX	10MEQ	A206630 001	Mar 29, 2019
	15MEQ	A206630 002	Mar 29, 2019
	20MEQ	A206630 003	Mar 29, 2019
	20MEQ	A210098 001	Apr 26, 2019
COPLEY PHARM	8MEQ	A070618 001	Sep 09, 1987
NESHER PHARMS	20MEQ	A076044 001	Apr 05, 2002
+ SCHERING	10MEQ **	N019439 002	Jun 13, 1986
+	20MEQ **	N019439 001	Jun 13, 1986
SIGMAPHARM LABS LLC	8MEQ	A207528 001	Aug 19, 2016
	10MEQ	A207528 002	Aug 19, 2016
STRIDES PHARMA	8MEQ	A206881 001	Jan 22, 2019
	10MEQ	A206881 002	Jan 22, 2019
	10MEQ	A210097 001	Jun 17, 2019
SLOW-K			
NOVARTIS	8MEQ	N017476 002	
TEN-K			
NOVARTIS	10MEQ	N019381 001	Apr 16, 1986

POTASSIUM CHLORIDE; SODIUM CHLORIDE

INJECTABLE; INJECTION

POTASSIUM CHLORIDE 0.037% IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER			
B BRAUN	37MG/100ML;900MG/100ML	N019708 001	Sep 29, 1989
POTASSIUM CHLORIDE 0.075% IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER			
B BRAUN	75MG/100ML;900MG/100ML	N019708 002	Sep 29, 1989
POTASSIUM CHLORIDE 0.11% IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER			
B BRAUN	110MG/100ML;900MG/100ML	N019708 003	Sep 29, 1989
POTASSIUM CHLORIDE 0.22% IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER			
B BRAUN	220MG/100ML;900MG/100ML	N019708 005	Sep 29, 1989
POTASSIUM CHLORIDE 0.224% IN SODIUM CHLORIDE 0.9%			
+ BAXTER HLTHCARE	224MG/100ML;900MG/100ML	N017648 003	
POTASSIUM CHLORIDE 0.3% IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER			
B BRAUN	300MG/100ML;900MG/100ML	N019708 006	Sep 29, 1989
SODIUM CHLORIDE 0.9% AND POTASSIUM CHLORIDE 0.075%			
BAXTER HLTHCARE	75MG/100ML;900MG/100ML	N017648 004	
SODIUM CHLORIDE 0.9% AND POTASSIUM CHLORIDE 0.075% IN PLASTIC CONTAINER			
B BRAUN	75MG/100ML;900MG/100ML	N018722 001	Nov 09, 1982
SODIUM CHLORIDE 0.9% AND POTASSIUM CHLORIDE 0.15% IN PLASTIC CONTAINER			
B BRAUN	150MG/100ML;900MG/100ML	N018722 002	Nov 09, 1982
SODIUM CHLORIDE 0.9% AND POTASSIUM CHLORIDE 0.22% IN PLASTIC CONTAINER			
B BRAUN	220MG/100ML;900MG/100ML	N018722 003	Nov 09, 1982
SODIUM CHLORIDE 0.9% AND POTASSIUM CHLORIDE 0.3% IN PLASTIC CONTAINER			
B BRAUN	300MG/100ML;900MG/100ML	N018722 004	Nov 09, 1982

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

POTASSIUM CHLORIDE; SODIUM CHLORIDE; TROMETHAMINE

INJECTABLE; INJECTION

THAM-E

HOSPIRA 370MG/VIAL; 1.75GM/VIAL; 36GM/VIAL N013025 001

POTASSIUM CITRATE

FOR SOLUTION; ORAL

POTASSIUM CITRATE

+ UT SW MEDCTR 10MEQ/PACKET ** N019647 002 Oct 13, 1988

+ 20MEQ/PACKET ** N019647 001 Oct 13, 1988

TABLET, EXTENDED RELEASE; ORAL

POTASSIUM CITRATE

TORRENT 5MEQ A213986 001 Feb 06, 2024

10MEQ A213986 002 Feb 06, 2024

15MEQ A213986 003 Feb 06, 2024

POTASSIUM IODIDE

SOLUTION; ORAL

POTASSIUM IODIDE

+ ROXANE 1GM/ML ** N018551 001 Feb 19, 1982

THYROSHIELD

ARCO PHARMS LLC 65MG/ML A077218 001 Jan 12, 2005

TABLET; ORAL

THYRO-BLOCK

MEDA PHARMS 130MG N018307 001

POTASSIUM PERCHLORATE

CAPSULE; ORAL

PERCHLORACAP

MALLINCKRODT 200MG N017551 001

POTASSIUM PHOSPHATE, DIBASIC; POTASSIUM PHOSPHATE, MONOBASIC

SOLUTION; INTRAVENOUS

POTASSIUM PHOSPHATES

+ CMP DEV LLC 4.5GM/15ML (300MG/ML); 2.65GM/15ML (175MG/ML) N212121 001 Sep 19, 2019

POVIDONE-IODINE

SOLUTION; TOPICAL

E-Z PREP

CLINIPAD 10% N019382 001 Jul 25, 1989

SPONGE; TOPICAL

E-Z PREP

CLINIPAD 5% N019382 002 Jul 25, 1989

E-Z PREP 220

CLINIPAD 5% N019382 003 Jul 25, 1989

E-Z SCRUB 201

+ BECTON DICKINSON 20% N019240 001 Nov 29, 1985

E-Z SCRUB 241

+ BECTON DICKINSON 10% N019476 001 Jan 07, 1987

PRALIDOXIME CHLORIDE

INJECTABLE; INJECTION

PRALIDOXIME CHLORIDE

BAXTER HLTHCARE CORP 300MG/ML N018799 001 Dec 13, 1982

SOLUTION; INTRAMUSCULAR

PRALIDOXIME CHLORIDE (AUTOINJECTOR)

+ MERIDIAN MEDCL TECHN 600MG/2ML (300MG/ML) N018986 001 Apr 26, 1983

TABLET; ORAL

PROTOPAM CHLORIDE

WYETH AYERST 500MG N014122 002

PRAMIPEXOLE DIHYDROCHLORIDE

TABLET; ORAL

MIRAPEX

+ BOEHRINGER INGELHEIM 0.125MG ** N020667 001 Jul 01, 1997

+ 0.25MG ** N020667 002 Jul 01, 1997

+ 0.5MG ** N020667 006 Feb 12, 1998

+ 0.75MG ** N020667 007 Jul 30, 2007

+ 1MG ** N020667 003 Jul 01, 1997

+ 1.25MG N020667 004 Jul 01, 1997

+ 1.5MG ** N020667 005 Jul 01, 1997

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PRAMIPEXOLE DIHYDROCHLORIDE

TABLET;ORAL

PRAMIPEXOLE DIHYDROCHLORIDE

ALEMBIC	0.125MG	A078894 001	Oct 08, 2010
	0.25MG	A078894 002	Oct 08, 2010
	0.5MG	A078894 003	Oct 08, 2010
	1MG	A078894 004	Oct 08, 2010
	1.5MG	A078894 005	Oct 08, 2010
CHARTWELL RX	0.125MG	A090241 001	Oct 08, 2010
	0.25MG	A090241 002	Oct 08, 2010
	0.5MG	A090241 003	Oct 08, 2010
	0.75MG	A090241 004	Oct 08, 2010
	1MG	A090241 005	Oct 08, 2010
	1.5MG	A090241 006	Oct 08, 2010
HERITAGE PHARMA AVET	0.125MG	A077724 001	Feb 19, 2008
	0.125MG	A078551 001	Oct 08, 2010
	0.125MG	A091254 001	Nov 30, 2010
	0.25MG	A077724 002	Feb 19, 2008
	0.25MG	A078551 002	Oct 08, 2010
	0.25MG	A091254 002	Nov 30, 2010
	0.5MG	A077724 003	Feb 19, 2008
	0.5MG	A078551 003	Oct 08, 2010
	0.5MG	A091254 003	Nov 30, 2010
	0.75MG	A091254 004	Nov 30, 2010
	1MG	A077724 004	Feb 19, 2008
	1MG	A078551 004	Oct 08, 2010
	1MG	A091254 005	Nov 30, 2010
	1.5MG	A077724 005	Feb 19, 2008
	1.5MG	A078551 005	Oct 08, 2010
	1.5MG	A091254 006	Nov 30, 2010
MACLEODS PHARMS LTD	0.125MG	A202164 001	Sep 20, 2012
	0.25MG	A202164 002	Sep 20, 2012
	0.5MG	A202164 003	Sep 20, 2012
	1MG	A202164 004	Sep 20, 2012
	1.5MG	A202164 005	Sep 20, 2012
NATCO	0.125MG	A077854 001	Oct 08, 2010
	0.25MG	A077854 002	Oct 08, 2010
	0.5MG	A077854 003	Oct 08, 2010
	1MG	A077854 004	Oct 08, 2010
	1.5MG	A077854 005	Oct 08, 2010
NATCO PHARMA	0.75MG	A090764 001	Apr 09, 2010
NOSTRUM LABS INC	0.125MG	A091450 001	Oct 08, 2010
	0.25MG	A091450 002	Oct 08, 2010
	0.5MG	A091450 003	Oct 08, 2010
	1MG	A091450 004	Oct 08, 2010
	1.5MG	A091450 005	Oct 08, 2010
SANDOZ	0.125MG	A090190 001	Jul 06, 2010
	0.25MG	A090190 002	Jul 06, 2010
	0.5MG	A090190 003	Jul 06, 2010
	0.75MG	A090190 006	Oct 08, 2010
	1MG	A090190 004	Jul 06, 2010
	1.5MG	A090190 005	Jul 06, 2010
SUN PHARM INDS INC	0.125MG	A091683 001	Mar 27, 2013
	0.25MG	A091683 002	Mar 27, 2013
	0.5MG	A091683 003	Mar 27, 2013
	0.75MG	A091683 004	Mar 27, 2013
	1MG	A091683 005	Mar 27, 2013
	1.5MG	A091683 006	Mar 27, 2013
UNICHEM	0.125MG	A207011 001	Dec 19, 2018
	0.25MG	A207011 002	Dec 19, 2018
	0.5MG	A207011 003	Dec 19, 2018
	0.75MG	A207011 004	Dec 19, 2018
	1MG	A207011 005	Dec 19, 2018
	1.5MG	A207011 006	Dec 19, 2018

TABLET, EXTENDED RELEASE;ORAL

MIRAPEX ER

+	BOEHRINGER INGELHEIM	0.375MG	N022421 001	Feb 19, 2010
+		0.75MG	N022421 002	Feb 19, 2010
+		1.5MG	N022421 003	Feb 19, 2010

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PRAMIPEXOLE DIHYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL

MIRAPEX ER

+	2.25MG	N022421 006	Jun 17, 2011
+	3MG	N022421 004	Feb 19, 2010
+	3.75MG	N022421 007	Jun 17, 2011
+	4.5MG	N022421 005	Feb 19, 2010

PRAMIPEXOLE DIHYDROCHLORIDE

ENDO OPERATIONS

0.375MG	A202206 001	Feb 06, 2014
0.75MG	A202206 002	Feb 06, 2014
1.5MG	A202206 003	Feb 06, 2014
2.25MG	A202206 004	Feb 06, 2014
3MG	A202206 005	Feb 06, 2014
3.75MG	A202206 006	Feb 06, 2014
4.5MG	A202206 007	Feb 06, 2014

PRAMLINTIDE ACETATE

INJECTABLE;SUBCUTANEOUS

SYMLIN

ASTRAZENECA AB

EQ 3MG BASE/5ML (EQ 600MCG BASE/ML) N021332 001 Mar 16, 2005

PRASUGREL HYDROCHLORIDE

TABLET;ORAL

PRASUGREL

CHARTWELL RX

EQ 5MG BASE A205790 001 Oct 16, 2017
EQ 10MG BASE A205790 002 Oct 16, 2017

DR REDDYS

EQ 5MG BASE A205926 001 Jul 07, 2020
EQ 10MG BASE A205926 002 Jul 07, 2020

LUPIN LTD

EQ 5MG BASE A205930 001 Jan 09, 2023
EQ 10MG BASE A205930 002 Jan 09, 2023PRAVASTATIN SODIUM

TABLET;ORAL

PRAVACHOL

+	BRISTOL MYERS SQUIBB	10MG **	N019898 002	Oct 31, 1991
+		20MG **	N019898 003	Oct 31, 1991
+		40MG **	N019898 004	Mar 22, 1993
+		80MG **	N019898 008	Dec 18, 2001

PRAVASTATIN SODIUM

HISUN PHARM HANGZHOU

20MG A206061 001 Nov 23, 2018
40MG A206061 002 Nov 23, 2018
80MG A206061 003 Nov 23, 2018

NORVIUM BIOSCIENCE

10MG A077013 001 Oct 23, 2006
10MG A079187 001 May 27, 2010
20MG A077013 002 Oct 23, 2006
20MG A079187 002 May 27, 2010
40MG A077013 003 Oct 23, 2006
40MG A079187 003 May 27, 2010
80MG A077013 004 Dec 28, 2007
80MG A079187 004 May 27, 2010

PLIVA HRVATSKA DOO

10MG A077730 001 Nov 21, 2006
20MG A077730 002 Nov 21, 2006
30MG A077730 003 Nov 21, 2006
40MG A077730 005 Nov 21, 2006

RANBAXY LABS LTD

10MG A076445 001 Apr 23, 2007
20MG A076445 002 Apr 23, 2007
40MG A076445 003 Apr 23, 2007
80MG A076445 004 Apr 23, 2007

ZYDUS PHARMS USA

10MG A077751 001 Apr 30, 2008

PRAZEPAM

CAPSULE;ORAL

CENTRAX

PARKE DAVIS

5MG N018144 001
10MG N018144 002
20MG N018144 003 May 10, 1982

PRAZEPAM

USL PHARMA

5MG A070427 001 Nov 06, 1987
10MG A070428 001 Nov 06, 1987

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PRAZEPAM

TABLET; ORAL

CENTRAX

PARKE DAVIS

10MG

N017415 001

PRAZQUANTEL

TABLET; ORAL

BILTRICIDE

+ BAYER HLTHCARE

600MG

N018714 001 Dec 29, 1982

PRAZOSIN HYDROCHLORIDE

CAPSULE; ORAL

PRAZOSIN HYDROCHLORIDE

AM THERAP

EQ 1MG BASE

A072782 001 May 16, 1989

EQ 2MG BASE

A072783 001 May 16, 1989

EQ 5MG BASE

A072784 001 May 16, 1989

ANI PHARMS

EQ 1MG BASE

A072577 002 May 16, 1989

EQ 2MG BASE

A072577 001 May 16, 1989

EQ 5MG BASE

A072577 003 May 16, 1989

DAVA PHARMS INC

EQ 1MG BASE

A072705 001 May 16, 1989

EQ 2MG BASE

A072706 001 May 16, 1989

EQ 5MG BASE

A072707 001 May 16, 1989

LANNETT CO INC

EQ 1MG BASE

A214083 001 Jan 03, 2024

EQ 2MG BASE

A214083 002 Jan 03, 2024

EQ 5MG BASE

A214083 003 Jan 03, 2024

PUREPAC PHARM

EQ 1MG BASE

A072991 001 May 16, 1989

EQ 2MG BASE

A072921 001 May 16, 1989

EQ 5MG BASE

A072992 001 May 16, 1989

WATSON LABS

EQ 1MG BASE

A072352 001 May 16, 1989

EQ 2MG BASE

A072333 001 May 16, 1989

EQ 5MG BASE

A072609 001 May 16, 1989

TABLET, EXTENDED RELEASE; ORAL

MINIPRESS XL

PFIZER

2.5MG

N019775 001 Jan 29, 1992

5MG

N019775 002 Jan 29, 1992

PREDNICARBATE

CREAM; TOPICAL

DERMATOP E EMOLLIENT

+ VALEANT BERMUDA

0.1% **

N020279 001 Oct 29, 1993

PREDNICARBATE

FOUGERA PHARMS

0.1%

A077287 001 Sep 19, 2006

OINTMENT; TOPICAL

DERMATOP

+ VALEANT PHARMS NORTH

0.1% **

N019568 001 Sep 23, 1991

PREDNISOLONE

CREAM; TOPICAL

METI-DERM

SCHERING

0.5%

N010209 002

SYRUP; ORAL

PREDNISOLONE

HIKMA

15MG/5ML

A040401 001 Feb 27, 2003

IVAX SUB TEVA PHARMS

15MG/5ML

A040287 001 May 28, 1999

NESHER PHARMS

5MG/5ML

A040423 001 Oct 22, 2001

15MG/5ML

A040364 001 Apr 10, 2002

PHARM ASSOC

5MG/5ML

A040570 001 Aug 25, 2005

15MG/5ML

A040399 001 Mar 05, 2003

PHARMOBEDIANT CNSLTG

15MG/5ML

A040313 001 Sep 10, 2003

TEVA PHARMS

15MG/5ML

A040322 001 Jan 19, 2000

WE PHARMS

15MG/5ML

A040192 001 May 28, 1998

PRELONE

MURO

5MG/5ML

A089654 001 Jan 17, 1989

TABLET; ORAL

CORTALONE

HALSEY

1MG

A080304 003

2.5MG

A080304 002

5MG

A080304 001

DELTA-CORTEF

PHARMACIA AND UPJOHN

5MG

N009987 004

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PREDNISOLONE

TABLET; ORAL

FERNISOLONE-P

FERNDALE LABS	5MG	A083941	001
---------------	-----	---------	-----

PREDNISOLONE

BARR	5MG	A084426	002
BUNDY	5MG	A083675	001
CHARTWELL RX	5MG	A084542	001
ELKINS SINN	5MG	A080625	001
EVERYLIFE	1MG	A084439	001
	2.5MG	A084439	002
	5MG	A084439	003
FERRANTE	2.5MG	A080562	001
	5MG	A080562	002
FOSUN PHARMA	5MG	A080339	001
+ HEATHER	5MG	A080326	001
IMPAX LABS	5MG	A080780	001
INWOOD LABS	5MG	A080748	001
IVAX SUB TEVA PHARMS	5MG	A080378	001
MARSHALL PHARMA	5MG	A080307	001
PANRAY	1MG	A080351	001
	5MG	A080351	002
PHOENIX LABS NY	5MG	A080322	001
PUREPAC PHARM	5MG	A080325	001
PVT FORM	5MG	A080211	001
RISING	5MG	A084773	001
ROXANE	5MG	A080327	002
SPERTI	1MG	A080358	001
	2.5MG	A080358	002
	5MG	A080358	003
SUPERPHARM	5MG	A088892	001 Feb 26, 1985
TABLICAPS	5MG	A085170	001
TEVA	5MG	A080398	001
UDL	5MG	A087987	001 Jan 18, 1983
VALEANT PHARM INTL	5MG	A080236	001
VITARINE	5MG	A080534	001
WATSON LABS	5MG	A085085	002
	5MG	A085415	001
	5MG	A085416	001
WEST WARD	5MG	A080324	001
WHITEWORTH TOWN PLSN	5MG	A080342	001

STERANE

PFIZER	5MG	N009996	001
--------	-----	---------	-----

PREDNISOLONE ACETATE

INJECTABLE; INJECTION

METICORTELONE

SCHERING	25MG/ML	N010255	002
----------	---------	---------	-----

PREDNISOLONE ACETATE

BEL MAR	25MG/ML	A083738	001
	50MG/ML	A083738	002
CENT PHARMS	25MG/ML	A084717	001
	50MG/ML	A084717	002
EPIC PHARMA LLC	25MG/ML	A083032	001
	50MG/ML	A084492	001
WATSON LABS	25MG/ML	A083398	001
	25MG/ML	A083654	001
	40MG/ML	A083767	001
	50MG/ML	A083764	001
	50MG/ML	A085781	001

STERANE

PFIZER	25MG/ML	N011446	001
--------	---------	---------	-----

SUSPENSION; ORAL

FLO-PRED

TARO	EQ 5MG BASE/5ML	N022067	001 Jan 17, 2008
	EQ 15MG BASE/5ML	N022067	002 Jan 17, 2008

SUSPENSION/DROPS; OPHTHALMIC

ECONOPRED

HARROW EYE	0.125%	N017468	001
------------	--------	---------	-----

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PREDNISOLONE ACETATE; SULFACETAMIDE SODIUM

OINTMENT;OPHTHALMIC

BLEPHAMIDE S.O.P.

ALLERGAN 0.2%;10% A087748 001 Dec 03, 1986

CETAPRED

ALCON 0.25%;10% A087771 001 Aug 06, 1993

METIMYD

SCHERING 0.5%;10% N010210 002 Sep 09, 1984

PREDSULFAIR

PHARMAFAIR 0.5%;10% A088032 001 Apr 15, 1983

VASOCIDIN

NOVARTIS 0.5%;10% A088791 001 Oct 05, 1984

SUSPENSION;OPHTHALMIC

BLEPHAMIDE

+ ALLERGAN 0.2%;10% N012813 002

ISOPTO CETAPRED

ALCON 0.25%;10% A087547 001

SUSPENSION/DROPS;OPHTHALMIC

METIMYD

SCHERING 0.5%;10% N010210 001

PREDAMIDE

EPIC PHARMA LLC 0.5%;10% A088059 001 Jul 29, 1983

PREDSULFAIR

PHARMAFAIR 0.5%;10% A088007 001 Apr 19, 1983

PREDSULFAIR II

PHARMAFAIR 0.2%;10% A088837 001 Dec 24, 1985

SULPHRIN

BAUSCH AND LOMB 0.5%;10% A088089 001 Dec 28, 1982

PREDNISOLONE SODIUM PHOSPHATE

INJECTABLE;INJECTION

HYDELTRASOL

MERCCK EQ 20MG PHOSPHATE/ML N011583 002

PREDNISOLONE SODIUM PHOSPHATE

WATSON LABS EQ 20MG PHOSPHATE/ML A080517 001

OINTMENT;OPHTHALMIC, OTIC

HYDELTRASOL

MERCCK EQ 0.25% PHOSPHATE N011028 001

SOLUTION;ORAL

ORAPRED

CONCORDIA PHARMS INC EQ 15MG BASE/5ML ** A075117 001 Dec 14, 2000

PREDNISOLONE SODIUM PHOSPHATE

AMNEAL PHARMS EQ 15MG BASE/5ML A078345 001 Mar 10, 2009

BAUSCH EQ 15MG BASE/5ML A075250 001 Jul 12, 2002

ENDO OPERATIONS EQ 15MG BASE/5ML A079010 001 May 26, 2009

HIKMA EQ 5MG BASE/5ML A075183 001 Mar 26, 2003

NESHER PHARMS EQ 5MG BASE/5ML A076982 001 May 24, 2005

EQ 15MG BASE/5ML A076988 001 May 24, 2005

PHARM ASSOC EQ 5MG BASE/5ML A076123 001 Dec 23, 2002

EQ 30MG BASE/5ML A204962 001 Mar 11, 2020

PHARMOBEDIENT EQ 15MG BASE/5ML A076895 001 Oct 04, 2004

PHARMOBEDIENT CNSLTG EQ 5MG BASE/5ML A075099 001 Jun 28, 2002

VINTAGE PHARMS EQ 5MG BASE/5ML A078416 001 Oct 31, 2007

WE PHARMS EQ 5MG BASE/5ML A075181 001 Dec 23, 2002

SOLUTION/DROPS;OPHTHALMIC

INFLAMASE FORTE

+ NOVARTIS EQ 0.9% PHOSPHATE A080751 002

INFLAMASE MILD

+ NOVARTIS EQ 0.11% PHOSPHATE A080751 001

METRETON

SCHERING EQ 0.5% PHOSPHATE A083834 001

PREDAIR

PHARMAFAIR EQ 0.11% PHOSPHATE A088415 001 Feb 29, 1984

PREDAIR FORTE

PHARMAFAIR EQ 0.9% PHOSPHATE A088165 001 Mar 28, 1983

PREDNISOLONE SODIUM PHOSPHATE

ALCON PHARMS LTD EQ 0.11% PHOSPHATE A081043 001 Oct 24, 1991

EQ 0.9% PHOSPHATE A081044 001 Oct 24, 1991

BAUSCH AND LOMB EQ 0.11% PHOSPHATE A040065 001 Jul 29, 1994

EPIC PHARMA LLC EQ 0.11% PHOSPHATE A083358 001

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PREDNISOLONE SODIUM PHOSPHATE

SOLUTION/DROPS;OPHTHALMIC

PREDNISOLONE SODIUM PHOSPHATE

	EQ 0.9% PHOSPHATE	A083358	002	
SOLA BARNES HIND	EQ 0.11% PHOSPHATE	A084171	001	
	EQ 0.9% PHOSPHATE	A084168	001	
	EQ 0.9% PHOSPHATE	A084169	001	
	EQ 0.9% PHOSPHATE	A084172	001	

TABLET, ORALLY DISINTEGRATING;ORAL

PREDNISOLONE SODIUM PHOSPHATE

RISING	EQ 10MG BASE	A202179	001	Apr 10, 2013
	EQ 15MG BASE	A202179	002	Apr 10, 2013
	EQ 30MG BASE	A202179	003	Apr 10, 2013

PREDNISOLONE SODIUM PHOSPHATE; SULFACETAMIDE SODIUM

SOLUTION/DROPS;OPHTHALMIC

SULFACETAMIDE SODIUM AND PREDNISOLONE SODIUM PHOSPHATE

GENUS	EQ 0.23% PHOSPHATE;10%	A073630	001	May 27, 1993
SULSTER				
EPIC PHARMA LLC	EQ 0.23% PHOSPHATE;10%	A074511	001	Jul 30, 1996
VASOCIDIN				
+ NOVARTIS	EQ 0.23% PHOSPHATE;10% **	N018988	001	Aug 26, 1988

PREDNISOLONE TEBUTATE

INJECTABLE;INJECTION

HYDELTRA-TBA

MERCK	20MG/ML	N010562	001	
-------	---------	---------	-----	--

PREDNISOLONE TEBUTATE

WATSON LABS	20MG/ML	A083362	001	Feb 17, 1984
-------------	---------	---------	-----	--------------

PREDNISON

SOLUTION;ORAL

PREDNISON

XTRIUM LABS INC	5MG/5ML	A089726	001	Aug 02, 1988
-----------------	---------	---------	-----	--------------

SYRUP;ORAL

LIQUID PRED

MURO	5MG/5ML	A087611	002	Sep 07, 1982
------	---------	---------	-----	--------------

TABLET;ORAL

CORTAN

HALSEY	20MG	A087480	001	
--------	------	---------	-----	--

DELTA-DOME

BAYER PHARMS	5MG	A080293	001	
--------------	-----	---------	-----	--

DELTASONE

+ PHARMACIA AND UPJOHN	2.5MG **	N009986	005	
	5MG **	N009986	002	
	10MG **	N009986	006	
	20MG **	N009986	007	
	50MG **	N009986	008	

FERNISON

FERNDALE LABS	5MG	A083364	001	
---------------	-----	---------	-----	--

METICORTEN

+ SCHERING	1MG **	N009766	002	
------------	--------	---------	-----	--

	5MG **	N009766	001	
--	--------	---------	-----	--

ORASONE

SOLVAY	1MG	A083009	001	
	5MG	A083009	002	
	10MG	A083009	003	
	20MG	A083009	004	
	50MG	A085999	001	

PARACORT

PARKE DAVIS	5MG	N010962	002	
-------------	-----	---------	-----	--

PREDNICEN-M

SCHWARZ PHARMA	5MG	A084655	001	
----------------	-----	---------	-----	--

PREDNISON

AM THERAP	5MG	A089387	001	Nov 06, 1986
	10MG	A089388	001	Nov 06, 1986
	20MG	A089389	001	Nov 06, 1986
AMNEAL PHARMS NY	5MG	A089597	001	Oct 05, 1987
	10MG	A089598	001	Oct 05, 1987
	20MG	A089599	001	Oct 05, 1987
BUNDY	5MG	A083676	001	

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PREDNISON

TABLET; ORAL

PREDNISON

CHARTWELL MOLECULAR	5MG	A080514	001	
CHARTWELL RX	5MG	A083059	001	
CONTRACT PHARMACAL	5MG	A080209	001	
DURAMED PHARMS BARR	5MG	A088394	001	Oct 04, 1983
	10MG	A088395	001	Oct 04, 1983
	20MG	A088396	001	Oct 04, 1983
ELKINS SINN	5MG	A080491	001	
	20MG	A085811	001	
EVERYLIFE	1MG	A084440	001	
	2.5MG	A084440	002	
	5MG	A084440	003	
FERRANTE	2.5MG	A080563	001	
	5MG	A080563	002	
GENEYORK PHARMS	1MG	A211496	001	Dec 28, 2018
	2.5MG	A210525	004	Dec 07, 2018
	5MG	A210525	005	Dec 07, 2018
	10MG	A210525	001	Dec 04, 2018
	20MG	A210525	002	Dec 04, 2018
	50MG	A210525	003	Dec 04, 2018
HALSEY	5MG	A080300	001	
HEATHER	5MG	A080320	001	
	10MG	A084341	001	
	20MG	A084417	001	
	20MG	A085543	001	
	50MG	A086946	001	
HIKMA PHARMS	1MG	A040890	001	Nov 01, 2010
	2.5MG	A040538	001	Jan 08, 2004
	50MG	A088465	001	Jun 01, 1984
IMPAX LABS	5MG	A080782	001	
INWOOD LABS	1MG	A080328	001	
	2.5MG	A080306	001	
	5MG	A080279	001	
IVAX SUB TEVA PHARMS	5MG	A080283	001	
	10MG	A084133	001	
	20MG	A084134	001	
KV PHARM	5MG	A084236	001	
LEDERLE	5MG	A086968	001	
MARSHALL PHARMA	5MG	A080301	001	
MUTUAL PHARM	5MG	A080701	001	
	10MG	A086595	001	
	20MG	A084634	001	
NYLOS	5MG	A085115	001	
PANRAY	1MG	A080350	001	
	2.5MG	A080350	002	
	5MG	A080350	003	
PHARMAVITE	5MG	A084662	002	
PHOENIX LABS NY	5MG	A080321	001	
	20MG	A083807	001	
PUREPAC PHARM	5MG	A080353	001	
	10MG	A086062	001	
	20MG	A086061	001	
PVT FORM	20MG	A085151	001	
REXALL	5MG	A080232	001	
RISING	5MG	A084774	001	
	10MG	A089983	001	Jan 12, 1989
	20MG	A085813	001	
	50MG	A089984	001	Jan 12, 1989
ROXANE	20MG	N017109	001	
+	25MG	A087833	001	May 04, 1982
SANDOZ	5MG	A080336	002	
SCHERER LABS	5MG	A080371	001	
SPERTI	1MG	A080359	001	
	2.5MG	A080359	002	
	5MG	A080359	003	
SUN PHARM INDUSTRIES	50MG	A086596	001	
SUPERPHARM	5MG	A088865	001	Oct 25, 1984

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PREDNISONTABLET;ORAL
PREDNISON

	10MG	A088866	001	Oct 25, 1984
	20MG	A088867	001	Oct 25, 1984
TEVA	5MG	A080397	001	
UDL	5MG	A087984	001	Jan 18, 1983
	10MG	A087985	001	Jan 18, 1983
	20MG	A087986	001	Jan 18, 1983
UPSHER SMITH	5MG	A087471	001	
	20MG	A087470	001	
VALEANT PHARM INTL	5MG	A080237	001	
VANGARD	5MG	A087682	001	Jan 15, 1982
	20MG	A087701	001	Jan 15, 1982
VITARINE	5MG	A080334	001	
	5MG	A080506	001	
WATSON LABS	5MG	A085084	002	
	10MG	A087773	001	Jul 13, 1982
	20MG	A086813	001	
	50MG	A086867	001	
	50MG	A087772	001	Jul 13, 1982
WHITEWORTH TOWN PLSN	2.5MG	A084913	001	
	5MG	A080343	001	
	10MG	A089028	001	Jul 24, 1986
	20MG	A084913	002	

SERVISONE

LEDERLE
TABLET, DELAYED RELEASE;ORAL
PREDNISON

	5MG	A080223	001	
ACTAVIS LABS FL INC	1MG	A204867	001	Apr 25, 2017
	2MG	A204867	002	Apr 25, 2017
	5MG	A204867	003	Apr 25, 2017
RAYOS				
+ HORIZON	1MG	N202020	001	Jul 26, 2012
+	2MG	N202020	002	Jul 26, 2012
+	5MG	N202020	003	Jul 26, 2012

PREGABALINCAPSULE;ORAL
PREGABALIN

CADILA PHARMS LTD	25MG	A206452	001	Jul 12, 2023
	50MG	A206452	002	Jul 12, 2023
	75MG	A206452	003	Jul 12, 2023
	100MG	A206452	004	Jul 12, 2023
	150MG	A206452	005	Jul 12, 2023
	200MG	A206452	006	Jul 12, 2023
	225MG	A206452	007	Jul 12, 2023
	300MG	A206452	008	Jul 12, 2023
CIPLA	25MG	A212280	001	Jan 10, 2020
	50MG	A212280	002	Jan 10, 2020
	75MG	A212280	003	Jan 10, 2020
	100MG	A212280	004	Jan 10, 2020
	150MG	A212280	005	Jan 10, 2020
	200MG	A212280	006	Jan 10, 2020
	225MG	A212280	007	Jan 10, 2020
	300MG	A212280	008	Jan 10, 2020
LUPIN LTD	25MG	A091040	001	May 03, 2022
	50MG	A091040	002	May 03, 2022
	75MG	A091040	003	May 03, 2022
	100MG	A091040	004	May 03, 2022
	150MG	A091040	005	May 03, 2022
	200MG	A091040	006	May 03, 2022
	225MG	A091040	007	May 03, 2022
	300MG	A091040	008	May 03, 2022
NORVIUM BIOSCIENCE	25MG	A091228	001	Sep 20, 2019
	50MG	A091228	002	Sep 20, 2019
	75MG	A091228	003	Sep 20, 2019
	100MG	A091228	004	Sep 20, 2019
	150MG	A091228	005	Sep 20, 2019
	200MG	A091228	006	Sep 20, 2019

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PREGABALINCAPSULE; ORAL
PREGABALIN

	225MG	A091228 007	Sep 20, 2019
	300MG	A091228 008	Sep 20, 2019
STRIDES PHARMA	25MG	A209883 001	Jan 24, 2024
	50MG	A209883 002	Jan 24, 2024
	75MG	A209883 003	Jan 24, 2024
	100MG	A209883 004	Jan 24, 2024
	150MG	A209883 005	Jan 24, 2024
	200MG	A209883 006	Jan 24, 2024
	225MG	A209883 007	Jan 24, 2024
	300MG	A209883 008	Jan 24, 2024
SUN PHARM	25MG	A091157 001	Nov 29, 2019
	50MG	A091157 002	Nov 29, 2019
	75MG	A091157 003	Nov 29, 2019
	100MG	A091157 004	Nov 29, 2019
	150MG	A091157 005	Nov 29, 2019
	200MG	A091157 006	Nov 29, 2019
	225MG	A091157 007	Nov 29, 2019
	300MG	A091157 008	Nov 29, 2019
TEVA PHARMS	25MG	A091219 001	Jul 19, 2019
	50MG	A091219 002	Jul 19, 2019
	75MG	A091224 001	Jul 19, 2019
	100MG	A091224 002	Jul 19, 2019
	150MG	A091224 003	Jul 19, 2019
	200MG	A091224 004	Jul 19, 2019
	225MG	A091224 005	Jul 19, 2019
	300MG	A091224 006	Jul 19, 2019
ZYDUS PHARMS	25MG	A206752 001	Dec 09, 2022
	50MG	A206752 002	Dec 09, 2022
	75MG	A206752 003	Dec 09, 2022
	100MG	A206752 004	Dec 09, 2022
	150MG	A206752 005	Dec 09, 2022
	200MG	A206752 006	Dec 09, 2022
	225MG	A206752 007	Dec 09, 2022
	300MG	A206752 008	Dec 09, 2022

TABLET, EXTENDED RELEASE; ORAL
PREGABALIN

ALVOGEN	82.5MG	A211687 001	Jul 06, 2021
	165MG	A211687 002	Jul 06, 2021
	330MG	A211687 003	Jul 06, 2021
NORVIUM BIOSCIENCE	82.5MG	A211948 001	Apr 13, 2021
	165MG	A211967 001	Nov 04, 2021
	330MG	A211431 001	Jul 02, 2021
SCIEGEN PHARMS INC	82.5MG	A215675 001	Sep 14, 2022
	165MG	A215675 002	Sep 14, 2022
	330MG	A215675 003	Sep 14, 2022
SUN PHARM	82.5MG	A211889 001	Apr 13, 2021
	165MG	A211889 002	Apr 13, 2021
	330MG	A211889 003	Apr 13, 2021
ZYDUS PHARMS	82.5MG	A215577 001	Aug 26, 2022
	165MG	A215577 002	Aug 26, 2022
	330MG	A215577 003	Aug 26, 2022

PRILOCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

CITANEST

+	ASTRAZENECA	1% **	N014763 004
+		2% **	N014763 005
+		3% **	N014763 003
	CITANEST PLAIN		
+	ASTRAZENECA	4% **	N014763 007
	CITANEST PLAIN DENTAL		
+	DENTSPLY PHARM	4%	N021382 001

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PRIMAQUINE PHOSPHATE

TABLET; ORAL

PRIMAQUINE PHOSPHATE

ALVOGEN

EQ 15MG BASE

A203924 001 Feb 03, 2014

PRIMIDONE

SUSPENSION; ORAL

MYSOLINE

+ FHTA

250MG/5ML

N010401 001

TABLET; ORAL

PRIMIDONE

DR REDDYS LABS LTD

50MG

A040862 001 Oct 03, 2008

250MG

A040862 002 Oct 03, 2008

HIKMA INTL PHARMS

50MG

A040667 001 Jul 27, 2006

250MG

A040667 002 Jul 27, 2006

IMPAX LABS

50MG

A040717 001 Feb 12, 2008

250MG

A040717 002 Feb 12, 2008

WATSON LABS

250MG

A085052 001

PROBENECID

TABLET; ORAL

BENEMID

+ MERCK

500MG **

N007898 004

PROBENECID

AUROBINDO PHARMA USA

500MG

A084211 002

IVAX SUB TEVA PHARMS

500MG

A083740 001 May 09, 1984

LEDERLE

500MG

A086917 001

WATSON LABS

500MG

A086150 002 Apr 23, 1982

PROBUCOL

TABLET; ORAL

LORELCO

SANOFI AVENTIS US

250MG

N017535 001

500MG

N017535 002 Jul 06, 1988

PROCAINAMIDE HYDROCHLORIDE

CAPSULE; ORAL

PROCAINAMIDE HYDROCHLORIDE

ANI PHARMS

250MG

A089219 001 Jul 01, 1986

375MG

A089219 002 Jul 01, 1986

500MG

A089219 003 Jul 01, 1986

ASCOT

250MG

A087542 001 Jan 08, 1982

375MG

A087697 001 Mar 01, 1983

500MG

A087543 001 Jan 08, 1982

IVAX SUB TEVA PHARMS

250MG

A084604 001

375MG

A084595 001

500MG

A084606 001

LANNETT

250MG

A083693 001

500MG

A084696 001

LEDERLE

250MG

A086942 001

375MG

A086952 001

500MG

A086943 001

ROXANE

250MG

A088989 001 Apr 26, 1985

500MG

A088990 001 Apr 26, 1985

VANGARD

250MG

A087643 001 Jun 01, 1982

500MG

A087875 001 Jun 01, 1982

WATSON LABS

250MG

A083287 001

250MG

A083795 001

250MG

A085167 001

375MG

A084403 001

375MG

A087020 001

500MG

A084280 001

500MG

A084357 001

500MG

A087021 001

PROCAN

PARKE DAVIS

250MG

A085804 001

375MG

A087502 001

500MG

A085079 001

PROCAPAN

PANRAY

250MG

A083553 002

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PROCAINAMIDE HYDROCHLORIDE

CAPSULE; ORAL

PRONESTYL

+	APOTHECON	250MG **	N007335 001
+		375MG **	N007335 004
+		500MG **	N007335 003

INJECTABLE; INJECTION

PROCAINAMIDE HYDROCHLORIDE

ABRAXIS PHARM	100MG/ML	A089415 001	Nov 17, 1986
	500MG/ML	A089416 001	Nov 17, 1986
HIKMA	100MG/ML	A089029 001	Apr 17, 1986
HOSPIRA	500MG/ML	A089537 001	Aug 25, 1987
INTL MEDICATION	500MG/ML	A088637 001	Jul 31, 1984
PHARMAFAIR	100MG/ML	A088824 001	Nov 20, 1985
	500MG/ML	A088830 001	Nov 20, 1985
SMITH AND NEPHEW	100MG/ML	A088530 001	Mar 04, 1985
	500MG/ML	A088531 001	Mar 04, 1985
SOLOPAK	500MG/ML	A088532 001	Mar 04, 1985
WARNER CHILCOTT	100MG/ML	A089528 001	May 03, 1988
	500MG/ML	A089529 001	May 03, 1988
WATSON LABS	100MG/ML	A087079 001	
	500MG/ML	A087080 001	
WEST-WARD PHARMS INT	500MG/ML	A089030 001	Apr 17, 1986

PRONESTYL

+	APOTHECON	100MG/ML **	N007335 002
+		500MG/ML **	N007335 005

TABLET; ORAL

PRONESTYL

APOTHECON	250MG	N017371 001
	375MG	N017371 002
	500MG	N017371 003

TABLET, EXTENDED RELEASE; ORAL

PROCAINAMIDE HYDROCHLORIDE

ANI PHARMS	250MG	A088958 001	Dec 02, 1985
	250MG	A089369 001	Aug 14, 1987
	500MG	A088959 001	Dec 02, 1985
	500MG	A088974 001	Jul 22, 1985
	500MG	A089369 002	Jan 09, 1987
	750MG	A089369 003	Aug 14, 1987
	750MG	A089438 001	Mar 23, 1987
	1GM	A040111 001	Dec 13, 1996
INWOOD LABS	500MG	A089840 001	Mar 06, 1989
SANDOZ	500MG	A089284 001	Jun 23, 1986
WATSON LABS	250MG	A088533 001	Dec 03, 1984
	250MG	A089026 001	Oct 22, 1985
	500MG	A088534 001	Dec 03, 1984
	500MG	A089027 001	Oct 22, 1985
	750MG	A088535 001	Nov 03, 1984
	750MG	A089042 001	Oct 22, 1985
	1GM	A089520 001	Jan 15, 1987

PROCAN SR

+	PARKE DAVIS	250MG	A086468 001	
+	PARKE DALE	500MG	A086065 001	
+		750MG	A087510 001	Apr 01, 1982
		1GM	A088489 001	Jan 16, 1985

PROCANBID

KING PHARMS	500MG	N020545 001	Jan 31, 1996
	1GM	N020545 002	Jan 31, 1996

PRONESTYL-SR

APOTHECON	500MG	A087361 001
-----------	-------	-------------

PROCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

NOVOCAIN

HOSPIRA	1%	A085362 003
	2%	A085362 004
	10%	A086797 001

PROCAINE HYDROCHLORIDE

ABRAXIS PHARM	1%	A080384 002
	1%	A080421 001

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PROCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

PROCAINE HYDROCHLORIDE

	2%	A080384	003
	2%	A080421	002
BEL MAR	1%	A080711	001
	2%	A080756	001
ELKINS SINN	1%	A083315	001
	2%	A083315	002
GD SEARLE LLC	1%	A086202	001
	2%	A086202	002
HOSPIRA	1%	A080416	001
	2%	A080416	002
MILES	1%	A080415	001
	2%	A080415	002
WATSON LABS	1%	A080658	001
	1%	A083535	001
	2%	A080658	002
	2%	A083535	002

PROCAINE HYDROCHLORIDE; TETRACYCLINE HYDROCHLORIDE

INJECTABLE; INJECTION

ACHROMYCIN

LEDERLE	40MG/VIAL; 100MG/VIAL	N050276	001
	40MG/VIAL; 250MG/VIAL	N050276	003

TETRACYN

PFIZER	40MG/VIAL; 100MG/VIAL	A060285	002
	40MG/VIAL; 250MG/VIAL	A060285	003

PROCAINE MERETHOXYLLINE; THEOPHYLLINE

INJECTABLE; INJECTION

DICURIN PROCAINE

LILLY	100MG/ML; 50MG/ML	N008869	001
-------	-------------------	---------	-----

PROCHLORPERAZINE

SUPPOSITORY; RECTAL

COMPAZINE

GLAXOSMITHKLINE	2.5MG **	N011127	003
	5MG **	N011127	001
	25MG **	N011127	002

PROCHLORPERAZINE

ABLE	2.5MG	A040407	001 Jul 11, 2001
	5MG	A040407	002 Jul 11, 2001
	25MG	A040407	003 Jul 11, 2001

PROCHLORPERAZINE EDISYLATE

CONCENTRATE; ORAL

COMPAZINE

+ GLAXOSMITHKLINE	EQ 10MG BASE/ML	N011276	001
-------------------	-----------------	---------	-----

PROCHLORPERAZINE

ALPHARMA US PHARMS	EQ 10MG BASE/ML	A087153	001 Jun 08, 1982
--------------------	-----------------	---------	------------------

PROCHLORPERAZINE EDISYLATE

MORTON GROVE	EQ 10MG BASE/ML	A088598	001 Oct 25, 1984
--------------	-----------------	---------	------------------

INJECTABLE; INJECTION

COMPAZINE

+ GLAXOSMITHKLINE	EQ 5MG BASE/ML **	N010742	002
-------------------	-------------------	---------	-----

PROCHLORPERAZINE

BAXTER HLTHCARE	EQ 5MG BASE/ML	A087759	001 Oct 01, 1982
-----------------	----------------	---------	------------------

PROCHLORPERAZINE EDISYLATE

HIKMA	EQ 5MG BASE/ML	A089523	001 May 03, 1988
-------	----------------	---------	------------------

	EQ 5MG BASE/ML	A213630	001 Nov 22, 2022
--	----------------	---------	------------------

HOSPIRA	EQ 5MG BASE/ML	A089703	001 Apr 07, 1988
---------	----------------	---------	------------------

MARSAM PHARMS LLC	EQ 5MG BASE/ML	A089675	001 Dec 05, 1988
-------------------	----------------	---------	------------------

SMITH AND NEPHEW	EQ 5MG BASE/ML	A089251	001 Dec 04, 1986
------------------	----------------	---------	------------------

TEVA PARENTERAL	EQ 5MG BASE/ML	A040505	001 May 30, 2003
-----------------	----------------	---------	------------------

WATSON LABS	EQ 5MG BASE/ML	A089530	001 Jul 08, 1987
-------------	----------------	---------	------------------

	EQ 5MG BASE/ML	A089605	001 Jul 08, 1987
--	----------------	---------	------------------

	EQ 5MG BASE/ML	A089606	001 Jul 08, 1987
--	----------------	---------	------------------

WYETH AYERST	EQ 5MG BASE/ML	A086348	001
--------------	----------------	---------	-----

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PROCHLORPERAZINE EDISYLATE

SYRUP;ORAL

COMPAZINE

GLAXOSMITHKLINE EQ 5MG BASE/5ML

N011188 001

PROCHLORPERAZINE EDISYLATE

ALPHARMA US PHARMS EQ 5MG BASE/5ML

A087154 001 Sep 01, 1982

MORTON GROVE EQ 5MG BASE/5ML

A088597 001 Oct 25, 1984

PROCHLORPERAZINE MALEATE

CAPSULE, EXTENDED RELEASE;ORAL

COMPAZINE

GLAXOSMITHKLINE EQ 10MG BASE

N011000 001

EQ 10MG BASE

N021019 001 Oct 06, 1999

EQ 15MG BASE

N011000 002

EQ 15MG BASE

N021019 002 Oct 06, 1999

EQ 30MG BASE

N011000 003

EQ 75MG BASE

N011000 004

TABLET;ORAL

COMPAZINE

+ GLAXOSMITHKLINE EQ 5MG BASE **

N010571 001

+ EQ 10MG BASE **

N010571 002

+ EQ 25MG BASE **

N010571 003

PROCHLORPERAZINE

WATSON LABS EQ 5MG BASE

A085580 001

EQ 10MG BASE

A085178 001

EQ 25MG BASE

A085579 001

PROCHLORPERAZINE MALEATE

CHARTWELL RX EQ 5MG BASE

A040101 001 Jul 19, 1996

EQ 10MG BASE

A040101 002 Jul 19, 1996

EQ 25MG BASE

A040101 003 Jul 19, 1996

DURAMED PHARMS BARR EQ 5MG BASE

A040207 001 May 01, 1997

EQ 5MG BASE

A089484 001 Jan 20, 1987

EQ 10MG BASE

A040207 002 May 01, 1997

EQ 10MG BASE

A089485 001 Jan 20, 1987

EQ 25MG BASE

A089486 001 Jan 20, 1987

IVAX SUB TEVA PHARMS EQ 5MG BASE

A040162 001 Jan 20, 1998

EQ 10MG BASE

A040162 002 Jan 20, 1998

NORVIUM BIOSCIENCE EQ 5MG BASE

A040185 002 Oct 28, 1996

EQ 10MG BASE

A040185 001 Oct 28, 1996

TEVA PHARMS EQ 5MG BASE

A040120 001 Jul 11, 1996

EQ 10MG BASE

A040120 002 Jul 11, 1996

PROCYCLIDINE HYDROCHLORIDE

TABLET;ORAL

KEMADRIN

MONARCH PHARMS 2MG

N009818 005

5MG

N009818 003

PROGESTERONE

CAPSULE;ORAL

PROGESTERONE

TEVA PHARMS 100MG

A202121 001 Feb 29, 2012

200MG

A202121 002 Feb 29, 2012

PROMETRIUM

VIRTUS 300MG

N019781 003 Oct 15, 1999

INJECTABLE; INJECTION

PROGESTERONE

ACCORD HLTHCARE 50MG/ML

A217707 001 Nov 21, 2023

+ ACTAVIS LABS UT INC 50MG/ML **

N017362 002

AM REGENT 50MG/ML

A090845 001 Jun 22, 2009

LILLY 25MG/ML

N009238 002

50MG/ML

N009238 001

INSERT, EXTENDED RELEASE; INTRAUTERINE

PROGESTASERT

ALZA 38MG

N017553 001

SYSTEM; VAGINAL

MILPROSA

+ FERRING PHARMS INC 1.78GM

N201110 001 Apr 29, 2020

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PROMAZINE HYDROCHLORIDE

CONCENTRATE; ORAL

SPARINE

WYETH AYERST	30MG/ML	N010942	001
	100MG/ML	N010942	004

INJECTABLE; INJECTION

PROMAZINE HYDROCHLORIDE

WATSON LABS	25MG/ML	A084510	001
	50MG/ML	A084517	001

SPARINE

HIKMA	25MG/ML	N010349	008
	50MG/ML	N010349	006

SYRUP; ORAL

SPARINE

WYETH AYERST	10MG/5ML	N010942	003
--------------	----------	---------	-----

TABLET; ORAL

SPARINE

WYETH AYERST	10MG	N010348	006
	25MG	N010348	001
	50MG	N010348	002
	100MG	N010348	003
	200MG	N010348	004

PROMETHAZINE HYDROCHLORIDE

INJECTABLE; INJECTION

PHENERGAN

WYETH AYERST	25MG/ML	N008857	002
	50MG/ML	N008857	003

PROMETHAZINE HYDROCHLORIDE

ABBOTT	25MG/ML	A084223	001
	50MG/ML	A084222	001
AM REGENT	25MG/ML	A040515	001 Mar 19, 2003
BEDFORD LABS	25MG/ML	A040524	001 Mar 17, 2004
	50MG/ML	A040524	002 Mar 17, 2004
EPIC PHARMA LLC	25MG/ML	A083955	002
	50MG/ML	A083955	001
HOSPIRA	25MG/ML	A040372	001 Jun 08, 2000
	50MG/ML	A040372	002 Jun 08, 2000
	50MG/ML	A083838	002
MARSAM PHARMS LLC	25MG/ML	A089463	001 May 02, 1988
	50MG/ML	A089477	001 May 02, 1988
MYLAN INSTITUTIONAL	25MG/ML	A040471	001 Nov 21, 2002
SANDOZ	25MG/ML	A040593	001 Nov 08, 2006
	50MG/ML	A040593	002 Nov 08, 2006
TEVA PHARMS USA	25MG/ML **	A040454	001 Aug 22, 2002
	50MG/ML **	A040454	002 Aug 22, 2002
WATSON LABS	25MG/ML	A083532	001
	25MG/ML	A084591	001
	50MG/ML	A080629	002
	50MG/ML	A083532	002
WOCKHARDT	25MG/ML	A040785	001 Sep 26, 2008
	50MG/ML	A040785	002 Sep 26, 2008

ZIPAN-25

ALTANA	25MG/ML	A083997	001
--------	---------	---------	-----

ZIPAN-50

ALTANA	50MG/ML	A083997	002
--------	---------	---------	-----

SUPPOSITORY; RECTAL

PHENERGAN

+ NORVIUM BIOSCIENCE	12.5MG **	N010926	002
+	25MG **	N010926	001
+	50MG **	N011689	001

PROMETHACON

POLYMEDICA	25MG	A084901	001
	50MG	A084902	001

PROMETHAZINE HYDROCHLORIDE

ABLE	12.5MG	A040504	001 Apr 11, 2003
	25MG	A040504	002 Apr 11, 2003
	50MG	A040449	001 Feb 27, 2003
WATSON LABS INC	12.5MG	A040479	001 Jun 24, 2003
	25MG	A040479	002 Jun 24, 2003

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PROMETHAZINE HYDROCHLORIDE

SYRUP;ORAL

MYMETHAZINE FORTIS				
USL PHARMA	25MG/5ML	A087996	001	Jan 18, 1983
PROMETH FORTIS				
ALPHARMA US PHARMS	25MG/5ML	A084772	001	
PROMETH PLAIN				
ACTAVIS MID ATLANTIC	6.25MG/5ML	A085953	001	
PROMETHAZINE				
CENCI	6.25MG/5ML	A089013	001	Sep 20, 1985
PROMETHAZINE HYDROCHLORIDE				
AMNEAL PHARMS	6.25MG/5ML	A040882	001	Dec 30, 2009
KV PHARM	6.25MG/5ML	A085388	001	
	25MG/5ML	A085385	001	
PHARM ASSOC	6.25MG/5ML	A040643	001	Apr 26, 2006
	6.25MG/5ML	A087518	001	
WHITEWORTH TOWN PLSN	6.25MG/5ML	A086395	001	
PROMETHAZINE HYDROCHLORIDE PLAIN				
+ ANI PHARMS	6.25MG/5ML **	N008381	004	Apr 18, 1984
+	25MG/5ML **	N008381	003	

TABLET;ORAL

PHENERGAN				
+ DELCOR ASSET CORP	12.5MG **	N007935	002	
+	25MG **	N007935	003	
+	50MG **	N007935	004	
PROMETHAZINE HYDROCHLORIDE				
ABBOTT	12.5MG	A084160	001	
	25MG	A084166	001	
	50MG	A084539	001	
ABLE	12.5MG	A040558	001	Jul 01, 2004
	25MG	A040558	002	Jul 01, 2004
	50MG	A040558	003	Jul 01, 2004
AUROBINDO PHARMA USA	12.5MG	A091054	001	Aug 30, 2011
	25MG	A091054	002	Aug 30, 2011
	50MG	A091054	003	Aug 30, 2011
CHARTWELL MOLECULAR	12.5MG	A080949	001	
	25MG	A080949	002	
	50MG	A080949	003	
IMPAX LABS	12.5MG	A040791	002	Feb 12, 2008
	25MG	A040791	003	Feb 12, 2008
	25MG	A084214	002	Jul 07, 1982
	50MG	A040791	001	May 20, 2008
INVATECH	12.5MG	A084233	001	
	25MG	A085146	001	
	50MG	A085146	002	
IVAX SUB TEVA PHARMS	12.5MG	A083604	001	
	25MG	A083603	001	
	50MG	A083613	001	
PVT FORM	12.5MG	A083214	001	
	25MG	A083658	001	
SANDOZ	12.5MG	A084176	002	May 22, 2009
SUN PHARM INDS INC	12.5MG	A040863	001	Dec 30, 2008
	25MG	A040863	002	Dec 30, 2008
	50MG	A040863	003	Dec 30, 2008
SUN PHARM INDUSTRIES	12.5MG	A084555	001	
	25MG	A084554	001	
	50MG	A084557	001	
TABLICAPS	12.5MG	A084080	001	
	25MG	A084027	001	
TEVA	25MG	A089109	001	Sep 10, 1985
WATSON LABS	12.5MG	A083401	001	
	12.5MG	A083712	001	
	12.5MG	A085986	001	
	25MG	A083204	001	
	25MG	A085684	001	
	50MG	A083403	001	
	50MG	A085664	001	
REMSSED				
BRISTOL MYERS SQUIBB	25MG	A083176	002	

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PROMETHAZINE HYDROCHLORIDETABLET; ORAL
REMSD

50MG A083176 001

PROPAFENONE HYDROCHLORIDECAPSULE, EXTENDED RELEASE; ORAL
PROPAFENONE HYDROCHLORIDE

MYLAN	225MG	A203803 001	Apr 29, 2016
	325MG	A203803 002	Apr 29, 2016
	425MG	A203803 003	Apr 29, 2016
TWI PHARMS	225MG	A212928 001	Jun 18, 2020
	325MG	A212928 002	Jun 18, 2020
	425MG	A212928 003	Jun 18, 2020

RYTHMOL SR

+ GLAXOSMITHKLINE LLC	225MG	N021416 001	Sep 04, 2003
+	325MG	N021416 002	Sep 04, 2003
+	425MG	N021416 003	Sep 04, 2003

TABLET; ORAL

PROPAFENONE HYDROCHLORIDE

NESHER PHARMS	150MG	A076193 001	Feb 07, 2002
	225MG	A076193 002	Feb 07, 2002
	300MG	A076193 003	Feb 07, 2002
SUN PHARM INDUSTRIES	150MG	A075998 001	Nov 29, 2001
	225MG	A075998 002	Nov 29, 2001
	300MG	A075998 003	Nov 29, 2001

RYTHMOL

+ GLAXOSMITHKLINE LLC	150MG **	N019151 001	Nov 27, 1989
+	225MG **	N019151 003	Nov 20, 1992
+	300MG **	N019151 002	Nov 27, 1989

PROPANTHELINE BROMIDE

INJECTABLE; INJECTION

PRO-BANTHINE

GD SEARLE LLC 30MG/VIAL N008843 001

TABLET; ORAL

PRO-BANTHINE

+ SHIRE	7.5MG **	N008732 003
+	15MG **	N008732 002

PROPANTHELINE BROMIDE

ASCOT	15MG	A087663 001	Oct 25, 1982
HEATHER	15MG	A085780 001	
HIKMA	7.5MG	A080927 001	
	15MG	A080927 002	
IMPAX LABS	15MG	A084541 002	
MYLAN	15MG	A083706 001	
PAR PHARM	15MG	A088377 001	Dec 08, 1983
PVT FORM	15MG	A080977 001	
SANDOZ	15MG	A080928 001	
TABLICAPS	15MG	A084428 001	
WATSON LABS	15MG	A083029 002	
	15MG	A083151 001	

PROPARACAINE HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

KAINAIR

PHARMAFAIR 0.5% A088087 001 Jun 07, 1983

OPHTHAINE

+ APOTHECON 0.5% ** N008883 001

OPHTHETIC

+ ALLERGAN 0.5% ** N012583 001

PARACAINE

OPTOPICS 0.5% A087681 001 Aug 05, 1982

PROPARACAINE HYDROCHLORIDE

SOLA BARNES HIND	0.5%	A084144 001
	0.5%	A084151 001

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PROPIOLACTONE

SOLUTION; IRRIGATION

BETAPRONE

FOREST LABS

N/A

N011657 001

PROPIOMAZINE HYDROCHLORIDE

INJECTABLE; INJECTION

LARGON

HIKMA

20MG/ML

N012382 002

PROPOFOL

INJECTABLE; INJECTION

DIPRIVAN

FRESENIUS KABI USA

10MG/ML

N019627 001 Oct 02, 1989

PROPOFOL

TEVA PARENTERAL

10MG/ML

A075392 001 Sep 19, 2000

WATSON LABS INC

10MG/ML

A205307 001 Dec 22, 2015

PROPOXYPHENE HYDROCHLORIDE

CAPSULE; ORAL

DARVON

XANODYNE PHARM

32MG

N010997 001

65MG

N010997 003

DOLENE

HERITAGE PHARMS INC

65MG

A080530 001

KESSO-GESIC

MK LABS

65MG

A083544 001

PROPHENE 65

HALSEY

65MG

A083538 002

PROPOXYPHENE HYDROCHLORIDE

ALRA

65MG

A083184 001

IMPAX LABS

65MG

A083317 001

IVAX SUB TEVA PHARMS

32MG

A083597 001

MUTUAL PHARM

65MG

A083186 001

MYLAN

32MG

A083528 001

65MG

A040569 001 Dec 16, 2004

65MG

A083299 001

NEXGEN PHARMA INC

65MG

A083185 001

PAR PHARM

65MG

A080269 001

PUREPAC PHARM

65MG

A083278 001

PVT FORM

32MG

A083464 001

65MG

A083113 001

ROXANE

32MG

A083089 001

65MG

A083089 002

SANDOZ

32MG

A084014 001

65MG

A083125 002

65MG

A083688 001

65MG

A083870 002

65MG

A086495 001

TEVA

65MG

A088615 001 Oct 22, 1984

VALEANT PHARM INTL

65MG

A080783 001

VINTAGE PHARMS

65MG

A040908 001 Jul 17, 2009

WATSON LABS

65MG

A080908 002

65MG

A085190 001

WEST WARD

65MG

A083501 001

WHITEWORTH TOWN PLSN

65MG

A084551 001

PROPOXYPHENE HYDROCHLORIDE 65

WARNER CHILCOTT

65MG

A083786 001

PROPOXYPHENE NAPSYLATE

SUSPENSION; ORAL

DARVON-N

AAIPHARMA LLC

50MG/5ML

N016861 001

TABLET; ORAL

DARVON-N

XANODYNE PHARM

100MG

N016862 002

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PROPRANOLOL HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE;ORAL

PROPRANOLOL HYDROCHLORIDE

EXTROVIS	60MG	A078022 001	Feb 15, 2007
	80MG	A078022 002	Feb 15, 2007
	120MG	A078022 003	Feb 15, 2007
	160MG	A078022 004	Feb 15, 2007
INWOOD LABS	60MG	A072499 001	Apr 11, 1989
	80MG	A072500 001	Apr 11, 1989
	120MG	A072501 001	Apr 11, 1989
	160MG	A072502 001	Apr 11, 1989
LUPIN LTD	60MG	A204349 001	Jan 12, 2024
	80MG	A204349 002	Jan 12, 2024
	120MG	A204349 003	Jan 12, 2024
	160MG	A204349 004	Jan 12, 2024
UPSHER SMITH LABS	60MG	A078311 001	Mar 06, 2009
	80MG	A078311 002	Mar 06, 2009
	120MG	A078311 003	Mar 06, 2009
	160MG	A078311 004	Mar 06, 2009

CONCENTRATE;ORAL

PROPRANOLOL HYDROCHLORIDE INTENSOL

ROXANE	80MG/ML	A071388 001	May 15, 1987
--------	---------	-------------	--------------

INJECTABLE; INJECTION

PROPRANOLOL HYDROCHLORIDE

+ BAXTER HLTHCARE CORP	1MG/ML **	N016419 001	
CHARTWELL INJECTABLE	1MG/ML	A075792 001	Aug 29, 2000
FOSUN PHARMA	1MG/ML	A076400 001	Feb 26, 2003
SMITH AND NEPHEW	1MG/ML	A070135 001	Apr 15, 1986
	1MG/ML	A070137 001	Apr 15, 1986
SOLOPAK	1MG/ML	A070136 001	Apr 15, 1986

SOLUTION;ORAL

PROPRANOLOL HYDROCHLORIDE

PAI HOLDINGS PHARM	20MG/5ML	A071984 001	Mar 03, 1989
	40MG/5ML	A071985 001	Mar 03, 1989

SUSPENSION;ORAL

INDERAL

WYETH AYERST	10MG/ML	N019536 001	Dec 12, 1986
--------------	---------	-------------	--------------

TABLET;ORAL

INDERAL

+ WYETH PHARMS	10MG **	N016418 001	
	20MG **	N016418 003	
	40MG **	N016418 002	
	60MG **	N016418 009	Oct 18, 1982
	80MG **	N016418 004	
	90MG **	N016418 010	Oct 18, 1982

PROPRANOLOL HYDROCHLORIDE

AIPING PHARM INC	90MG	A071288 001	Oct 22, 1986
ANI PHARMS	60MG	A071791 001	Jul 15, 1987
	90MG	A071977 001	Apr 06, 1988
CHARTWELL RX	10MG	A070663 001	Jun 13, 1986
	20MG	A070664 001	Jun 13, 1986
	40MG	A070665 001	Jun 13, 1986
	60MG	A070666 001	Oct 10, 1986
	80MG	A070667 001	Jun 13, 1986
DAVA PHARMS INC	10MG	A070125 001	Jul 30, 1985
	20MG	A070126 001	Jul 30, 1985
	40MG	A070127 001	Jul 30, 1985
	60MG	A071495 001	Dec 31, 1987
	80MG	A070128 001	Jul 30, 1985
	90MG	A071496 001	Dec 31, 1987
DURAMED PHARMS BARR	10MG	A070306 001	Sep 09, 1985
	20MG	A070307 001	Sep 09, 1985
	40MG	A070308 001	Sep 09, 1985
	60MG	A070309 001	Oct 01, 1986
	80MG	A070310 001	Sep 09, 1985
	90MG	A071327 001	Oct 01, 1986
INTERPHARM	10MG	A071368 001	May 05, 1987
	20MG	A071369 001	May 05, 1987
	40MG	A071370 001	May 05, 1987

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PROPRANOLOL HYDROCHLORIDE

TABLET; ORAL

PROPRANOLOL HYDROCHLORIDE

	80MG	A071371 001	May 05, 1987
IVAX SUB TEVA PHARMS	10MG	A072063 001	Jul 29, 1988
	20MG	A072066 001	Jul 29, 1988
	40MG	A072067 001	Jul 29, 1988
	60MG	A072068 001	Jul 29, 1988
	80MG	A072069 001	Jul 29, 1988
LEDERLE	10MG	A072117 001	Jun 23, 1988
	20MG	A072118 001	Jun 23, 1988
	40MG	A072119 001	Jun 23, 1988
	80MG	A072120 001	Jun 23, 1988
MYLAN	60MG	A072275 001	Jun 09, 1989
PUREPAC PHARM	10MG	A070814 001	Nov 03, 1986
	20MG	A070815 001	Nov 03, 1986
	40MG	A070816 001	Nov 03, 1986
	60MG	A070817 001	Nov 03, 1986
	80MG	A070757 001	Nov 03, 1986
ROXANE	10MG	A070516 001	Jul 07, 1986
	20MG	A070517 001	Jul 07, 1986
	40MG	A070518 001	Jul 07, 1986
	60MG	A070519 001	Sep 24, 1986
	80MG	A070520 001	Jul 07, 1986
	90MG	A070521 001	Sep 24, 1986
SANDOZ	10MG	A071658 001	Jul 05, 1988
	20MG	A071687 001	Jul 05, 1988
	40MG	A071688 001	Jul 05, 1988
	60MG	A072197 001	Jul 05, 1988
	80MG	A071689 001	Jul 05, 1988
	90MG	A072198 001	Jul 05, 1988
SCHERING	10MG	A070120 001	Aug 06, 1985
	20MG	A070121 001	Aug 06, 1985
	40MG	A070122 001	Aug 06, 1985
	60MG	A070123 001	Oct 29, 1986
	80MG	A070124 001	Aug 06, 1985
SUPERPHARM	10MG	A071515 001	Jun 08, 1988
	20MG	A071516 001	Jun 08, 1988
	40MG	A071517 001	Jun 08, 1988
	80MG	A071518 001	Jun 08, 1988
TEVA	10MG	A070232 001	Oct 07, 1987
	20MG	A070233 001	Jun 23, 1986
	40MG	A070234 001	Jun 23, 1986
WARNER CHILCOTT	10MG	A070438 001	Sep 15, 1986
	20MG	A070439 001	Sep 15, 1986
	40MG	A070440 001	Sep 15, 1986
	60MG	A070441 001	Sep 24, 1986
	80MG	A070442 001	Sep 15, 1986
WATSON LABS	10MG	A070140 001	Jul 30, 1985
	10MG	A070378 001	Mar 19, 1987
	20MG	A070141 001	Jul 30, 1985
	20MG	A070379 001	Mar 19, 1987
	40MG	A070142 001	Jul 30, 1985
	40MG	A070380 001	Mar 19, 1987
	60MG	A070143 001	Jan 15, 1987
	60MG	A070178 002	Apr 23, 2018
	60MG	A070381 001	Mar 19, 1987
	60MG	A071098 001	Oct 06, 1986
	80MG	A070144 001	Jul 30, 1985
	80MG	A070382 001	Mar 19, 1987
	80MG	A070551 001	Jul 10, 1986
	90MG	A071183 001	Oct 06, 1986
	90MG	A071792 001	Jul 15, 1987
WATSON LABS TEVA	10MG	A070548 001	Jul 10, 1986
	20MG	A070549 001	Apr 11, 1986
	40MG	A070550 001	Apr 11, 1986

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PROPYLIODONE

SUSPENSION; INTRATRACHEAL

DIONOSIL AQUEOUS

GLAXOSMITHKLINE

50%

N009309 001

DIONOSIL OILY

GLAXOSMITHKLINE

60%

N009309 002

PROPYLTHIOURACIL

TABLET; ORAL

PROPYLTHIOURACIL

ABBOTT

50MG

A084075 001

ANABOLIC

50MG

A080285 001

ANI PHARMS

50MG

A080215 001

CHARTWELL MOLECULAR

50MG

A080016 001

CHARTWELL RX

50MG

A084543 001

HALSEY

50MG

A080015 001

IMPAX LABS

50MG

A080159 001

LILLY

50MG

N006213 001

SUN PHARM INDUSTRIES

50MG

A083982 001

TABLICAPS

50MG

A080840 001

WATSON LABS

50MG

A080932 001

50MG

A085201 001

PROTAMINE SULFATE

INJECTABLE; INJECTION

PROTAMINE SULFATE

HIKMA

10MG/ML

A089474 001 Nov 05, 1986

10MG/ML

A089475 001 Nov 05, 1986

+ LILLY

10MG/ML **

N006460 002

PHARMACIA AND UPJOHN

50MG/VIAL

N007413 001

250MG/VIAL

N007413 002 Aug 02, 1984

PROTEIN HYDROLYSATE

INJECTABLE; INJECTION

AMINOSOL 5%

ABBVIE

5%

N005932 012 Jan 31, 1985

HYPROTIGEN 5%

B BRAUN

5%

N006170 003 Jan 10, 1984

PROTIRELIN

INJECTABLE; INJECTION

THYPINONE

ABBOTT

0.5MG/ML

N017638 001

THYREL TRH

FERRING

0.5MG/ML

N018087 001

PROTOKYLOL HYDROCHLORIDE

TABLET; ORAL

VENTAIRE

SANOFI AVENTIS US

2MG

A083459 001

PROTRIPTYLINE HYDROCHLORIDE

TABLET; ORAL

VIVACTIL

CHARTWELL RX

5MG

A073644 001 Aug 24, 1995

10MG

A073645 001 Aug 24, 1995

TEVA WOMENS

5MG **

N016012 001

10MG **

N016012 002

PSEUDOEPHEDRINE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

NOVAFED

SANOFI AVENTIS US

120MG

N017603 001

SUDAFED 12 HOUR

+ GLAXOSMITHKLINE

120MG **

N017941 002

TABLET, EXTENDED RELEASE; ORAL

SUDAFED 12 HOUR

MCNEIL CONS

120MG

A073585 001 Oct 31, 1991

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PSEUDOEPHEDRINE HYDROCHLORIDE; TRIPROLIDINE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE;ORAL

ACTIFED

GLAXOSMITHKLINE 120MG;5MG N018996 001 Jun 17, 1985

TRIPROLIDINE AND PSEUDOEPHEDRINE HYDROCHLORIDES

KV PHARM 120MG;5MG A071798 001 Mar 16, 1989

SYRUP;ORAL

ACTAHIST

CENCI 30MG/5ML;1.25MG/5ML A088344 001 Feb 09, 1984

HISTAFED

CENCI 30MG/5ML;1.25MG/5ML A088283 001 Apr 20, 1984

MYFED

USL PHARMA 30MG/5ML;1.25MG/5ML A088116 001 Mar 04, 1983

TRILITRON

NEWTRON PHARMS 30MG/5ML;1.25MG/5ML A088474 001 Feb 12, 1985

TABLET;ORAL

ALLERFED

PVT FORM 60MG;2.5MG A088860 001 Jan 31, 1985

CORPHED

CHARTWELL RX 60MG;2.5MG A088602 001 Apr 11, 1985

PSEUDOEPHEDRINE HYDROCHLORIDE AND TRIPROLIDINE HYDROCHLORIDE

SANDOZ 60MG;2.5MG A088193 001 May 17, 1983

TRILITRON

NEWTRON PHARMS 60MG;2.5MG A088515 001 Jan 09, 1985

TRIPHED

TEVA 60MG;2.5MG A088630 001 May 17, 1984

TRIPROLIDINE AND PSEUDOEPHEDRINE

WATSON LABS 60MG;2.5MG A088318 002 Jan 13, 1984

WEST WARD 60MG;2.5MG A088117 001 Apr 19, 1983

TRIPROLIDINE HYDROCHLORIDE AND PSEUDOEPHEDRINE HYDROCHLORIDE

IVAX SUB TEVA PHARMS 60MG;2.5MG A085273 001 Dec 12, 1984

SUPERPHARM 60MG;2.5MG A088578 001 Feb 21, 1985

TABLET, EXTENDED RELEASE;ORAL

TRIPROLIDINE AND PSEUDOEPHEDRINE HYDROCHLORIDES

KV PHARM 120MG;5MG A072758 001 Nov 25, 1991

PSEUDOEPHEDRINE POLISTIREX

SUSPENSION, EXTENDED RELEASE;ORAL

PSEUDO-12

UCB INC EQ 60MG HYDROCHLORIDE/5ML N019401 001 Jun 19, 1987

PSEUDOEPHEDRINE SULFATE

TABLET, EXTENDED RELEASE;ORAL

AFRINOL

+ SCHERING PLOUGH 120MG N018191 001

PYRIDOSTIGMINE BROMIDE

TABLET;ORAL

PYRIDOSTIGMINE BROMIDE

ANI PHARMS 30MG A040512 002 Jul 20, 2005

IMPAX LABS INC 60MG A040457 001 Dec 26, 2002

SOLVAY 30MG A089572 001 Nov 27, 1990

US ARMY 30MG N020414 001 Feb 05, 2003

TABLET, EXTENDED RELEASE;ORAL

PYRIDOSTIGMINE BROMIDE

AMNEAL 105MG N217604 001 Oct 04, 2024

PYRIDOXINE HYDROCHLORIDE

INJECTABLE;INJECTION

HEXA-BETALIN

LILLY 100MG/ML A080854 001

PYRIDOXINE HYDROCHLORIDE

BEL MAR 100MG/ML A080761 001

DELL LABS 50MG/ML A083771 001

100MG/ML A083772 001

DR REDDYS 100MG/ML A080572 001

ELKINS SINN 100MG/ML A080581 001

EPIC PHARMA LLC 100MG/ML A087967 001 Oct 01, 1982

LUITPOLD 100MG/ML A080669 001

MYLAN INSTITUTIONAL 100MG/ML A204879 001 Jul 14, 2016

WATSON LABS 100MG/ML A083760 001

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PYRILAMINE MALEATE

TABLET;ORAL

PYRILAMINE MALEATE

IMPAX LABS

25MG

A080808 001

WATSON LABS

25MG

A085231 001

PYRIMETHAMINE; SULFADOXINE

TABLET;ORAL

FANSIDAR

ROCHE

25MG;500MG

N018557 001

PYRITHIONE ZINC

LOTION;TOPICAL

HEAD & SHOULDERS CONDITIONER

WARNER CHILCOTT

0.3%

N019412 002 Mar 10, 1986

PYRVINIUM PAMOATE

SUSPENSION;ORAL

POVAN

PARKE DAVIS

EQ 50MG BASE/5ML

N011964 001

TABLET;ORAL

POVAN

PARKE DAVIS

EQ 50MG BASE

N012485 002

QUAZEPAM

TABLET;ORAL

DORAL

GALT PHARMS

7.5MG

N018708 003 Feb 26, 1987

QUETIAPINE FUMARATE

TABLET;ORAL

QUETIAPINE FUMARATE

ACTAVIS GRP PTC

EQ 25MG BASE

A201762 001 Feb 27, 2013

EQ 50MG BASE

A201762 002 Feb 27, 2013

EQ 100MG BASE

A201762 003 Feb 27, 2013

EQ 150MG BASE

A201762 004 Feb 27, 2013

EQ 200MG BASE

A201762 005 Feb 27, 2013

EQ 300MG BASE

A201762 006 Feb 27, 2013

EQ 400MG BASE

A201762 007 Feb 27, 2013

ALEMBIC

EQ 25MG BASE

A203390 001 Oct 28, 2014

EQ 50MG BASE

A203390 002 Oct 28, 2014

EQ 100MG BASE

A203390 003 Oct 28, 2014

EQ 200MG BASE

A203390 004 Oct 28, 2014

EQ 300MG BASE

A203390 005 Oct 28, 2014

EQ 400MG BASE

A203390 006 Oct 28, 2014

CHARTWELL RX

EQ 25MG BASE

A078679 001 Dec 14, 2012

EQ 50MG BASE

A078679 002 Dec 14, 2012

EQ 100MG BASE

A078679 003 Dec 14, 2012

EQ 150MG BASE

A078679 004 Dec 14, 2012

EQ 200MG BASE

A078679 005 Dec 14, 2012

EQ 300MG BASE

A078679 006 Dec 14, 2012

EQ 400MG BASE

A078679 007 Dec 14, 2012

IPCA LABS LTD

EQ 25MG BASE

A205983 001 May 20, 2024

EQ 50MG BASE

A205983 002 May 20, 2024

EQ 100MG BASE

A205983 003 May 20, 2024

EQ 200MG BASE

A205983 004 May 20, 2024

EQ 300MG BASE

A205983 005 May 20, 2024

EQ 400MG BASE

A205983 006 May 20, 2024

JUBILANT GENERICS

EQ 25MG BASE

A203150 001 Nov 26, 2013

NORVIUM BIOSCIENCE

EQ 25MG BASE

A090323 001 Mar 27, 2012

TORRENT PHARMS LTD

EQ 25MG BASE

A200363 001 Mar 27, 2012

EQ 50MG BASE

A200363 002 Mar 27, 2012

EQ 100MG BASE

A200363 003 Mar 27, 2012

EQ 200MG BASE

A200363 004 Mar 27, 2012

EQ 300MG BASE

A200363 005 Mar 27, 2012

EQ 400MG BASE

A200363 006 Mar 27, 2012

SEROQUEL

+ ASTRAZENECA

EQ 150MG BASE **

N020639 004 Dec 20, 1998

TABLET, EXTENDED RELEASE;ORAL

QUETIAPINE FUMARATE

AMNEAL PHARMS

EQ 400MG BASE

A211405 001 Oct 26, 2018

ENDO OPERATIONS

EQ 50MG BASE

A090482 001 May 09, 2017

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

QUETIAPINE FUMARATETABLET, EXTENDED RELEASE;ORAL
QUETIAPINE FUMARATE

	EQ 150MG BASE	A090482 002	May 09, 2017
	EQ 150MG BASE	A090757 001	Dec 01, 2017
	EQ 200MG BASE	A090482 003	May 09, 2017
	EQ 200MG BASE	A090757 002	Dec 01, 2017
	EQ 300MG BASE	A090482 004	May 09, 2017
	EQ 300MG BASE	A090757 003	Dec 01, 2017
	EQ 400MG BASE	A090482 005	May 09, 2017
	EQ 400MG BASE	A090757 004	Dec 01, 2017
PHARMADAX INC	EQ 50MG BASE	A206260 001	May 09, 2017
	EQ 150MG BASE	A206260 002	May 09, 2017
	EQ 200MG BASE	A206260 003	May 09, 2017
	EQ 300MG BASE	A206260 004	May 09, 2017
	EQ 400MG BASE	A206260 005	May 09, 2017
RISING	EQ 50MG BASE	A202228 001	Feb 02, 2021
	EQ 150MG BASE	A202228 002	Feb 02, 2021
	EQ 200MG BASE	A202228 003	Feb 02, 2021
	EQ 300MG BASE	A202228 004	Feb 02, 2021
	EQ 400MG BASE	A202228 005	Feb 02, 2021

QUINAPRIL HYDROCHLORIDE

TABLET;ORAL

ACCUPRIL

+	PFIZER PHARMS	EQ 5MG BASE **	N019885 001	Nov 19, 1991
+		EQ 10MG BASE **	N019885 002	Nov 19, 1991
+		EQ 20MG BASE **	N019885 003	Nov 19, 1991
+		EQ 40MG BASE **	N019885 004	Nov 19, 1991

QUINAPRIL HYDROCHLORIDE

ACTAVIS ELIZABETH

	EQ 5MG BASE	A076459 001	Dec 22, 2004
	EQ 10MG BASE	A076459 002	Dec 22, 2004
	EQ 20MG BASE	A076459 003	Dec 22, 2004
	EQ 40MG BASE	A076459 004	Dec 22, 2004
ACTAVIS LABS FL INC	EQ 5MG BASE	A076049 001	Jan 14, 2005
	EQ 10MG BASE	A076049 002	Jan 14, 2005
	EQ 20MG BASE	A076049 003	Jan 14, 2005
	EQ 40MG BASE	A076049 004	Jan 14, 2005

ANI PHARMS

	EQ 5MG BASE	A075504 001	Aug 24, 2007
	EQ 10MG BASE	A075504 002	Aug 24, 2007
	EQ 20MG BASE	A075504 003	Aug 24, 2007
	EQ 40MG BASE	A075504 004	Aug 24, 2007

APOTEX INC

	EQ 5MG BASE	A076240 001	Jan 26, 2006
	EQ 10MG BASE	A076240 002	Jan 26, 2006
	EQ 20MG BASE	A076240 003	Jan 26, 2006
	EQ 40MG BASE	A076240 004	Jan 26, 2006

INVAGEN PHARMS

	EQ 5MG BASE	A078457 001	Aug 24, 2007
	EQ 10MG BASE	A078457 002	Aug 24, 2007
	EQ 20MG BASE	A078457 003	Aug 24, 2007
	EQ 40MG BASE	A078457 004	Aug 24, 2007

MYLAN

	EQ 5MG BASE	A076036 001	Jan 28, 2005
	EQ 10MG BASE	A076036 002	Jan 28, 2005
	EQ 20MG BASE	A076036 003	Jan 28, 2005
	EQ 40MG BASE	A076036 004	Jan 28, 2005

NORVIUM BIOSCIENCE

	EQ 5MG BASE	A076694 001	Dec 23, 2004
	EQ 10MG BASE	A076694 002	Dec 23, 2004
	EQ 20MG BASE	A076694 003	Dec 23, 2004
	EQ 40MG BASE	A076694 004	Dec 23, 2004

SUN PHARM INDS LTD

	EQ 5MG BASE	A076607 001	Dec 15, 2004
	EQ 5MG BASE	A090800 001	Jun 18, 2009
	EQ 10MG BASE	A076607 002	Dec 15, 2004
	EQ 10MG BASE	A090800 002	Jun 18, 2009
	EQ 20MG BASE	A076607 003	Dec 15, 2004
	EQ 20MG BASE	A090800 003	Jun 18, 2009
	EQ 40MG BASE	A076607 004	Dec 15, 2004
	EQ 40MG BASE	A090800 004	Jun 18, 2009

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

QUINESTROL

TABLET; ORAL

ESTROVIS

PARKE DAVIS

0.1MG

N016768 002

0.2MG

N016768 003

QUINETHAZONE

TABLET; ORAL

HYDROMOX

LEDERLE

50MG

N013264 001

QUINETHAZONE; RESERPINE

TABLET; ORAL

HYDROMOX R

LEDERLE

50MG; 0.125MG

N013927 001

QUINIDINE GLUCONATE

INJECTABLE; INJECTION

QUINIDINE GLUCONATE

+ LILLY

80MG/ML

N007529 002 Feb 10, 1989

TABLET; ORAL

QUINACT

BAYER HLTHCARE

266MG

A085978 001

400MG

A086099 001

TABLET, EXTENDED RELEASE; ORAL

DURAQUIN

WARNER CHILCOTT

330MG

N017917 001

QUINAGLUTE

+ BAYER HLTHCARE

324MG **

N016647 001

QUINALAN

CHARTWELL MOLECULAR

324MG

A088081 001 Feb 10, 1986

QUINATIME

WATSON LABS

324MG

A087448 001

QUINIDINE GLUCONATE

ANI PHARMS

324MG

A087810 001 Sep 29, 1982

ASCOT

324MG

A088582 001 Jun 17, 1985

CYCLE

324MG

A088431 001 Jan 06, 1984

HALSEY

324MG

A089476 001 Apr 10, 1987

RISING

324MG

A089894 001 Dec 15, 1988

SUPERPHARM

324MG

A089164 001 Nov 21, 1985

WATSON LABS

324MG

A087785 001 Jan 24, 1983

QUINIDINE POLYGALACTURONATE

TABLET; ORAL

CARDIOQUIN

PHARM RES ASSOC

275MG

N011642 002

QUINIDINE SULFATE

CAPSULE; ORAL

CIN-QUIN

SOLVAY

200MG

A085296 001

300MG

A085297 001

QUINIDINE SULFATE

LILLY

200MG

A085103 001

TABLET; ORAL

CIN-QUIN

+ SOLVAY

100MG

A085299 001

+

200MG

A084932 001

300MG

A085298 001

QUINIDINE SULFATE

BARR

200MG

A084177 001

CHARTWELL MOLECULAR

200MG

A083743 001

CHARTWELL RX

300MG

A089839 001 Sep 29, 1988

CONTRACT PHARMACAL

200MG

A083808 001

CYCLE

200MG

A083640 001

300MG

A085632 001

DAVA PHARMS INC

200MG

A087011 001

ELKINS SINN

200MG

A083622 001

EVERYLIFE

200MG

A083439 001

HALSEY

200MG

A083583 001

HIKMA

200MG

A083862 001

IMPAX LABS

200MG

A083347 001

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

QUINIDINE SULFATE

TABLET;ORAL

QUINIDINE SULFATE

IVAX SUB TEVA PHARMS	200MG	A084549	001	
KING PHARMS	200MG	A085175	001	
KV PHARM	200MG	A085276	001	
LEDERLE	200MG	A086176	001	
LILLY	200MG	A085038	001	
PERRIGO	200MG	A085322	001	
PHARMAVITE	200MG	A084627	001	
PUREPAC PHARM	200MG	A084003	001	
SANDOZ	200MG	A084631	001	
	200MG	A084914	001	
SCHERER LABS	200MG	A085068	001	
SUN PHARM INDUSTRIES	100MG	A081029	001	Apr 14, 1989
	200MG	A081030	001	Apr 14, 1989
	300MG	A081031	001	Apr 14, 1989
SUPERPHARM	200MG	A088973	001	Apr 10, 1985
USL PHARMA	200MG	A087837	001	Apr 14, 1982
VALEANT PHARM INTL	200MG	A083393	001	
VANGARD	200MG	A087909	001	Jul 13, 1982
VINTAGE PHARMS	200MG	A083963	001	
WARNER CHILCOTT	200MG	A083879	001	
WATSON LABS	100MG	A085584	001	
	200MG	A083288	001	
	200MG	A085140	002	
	300MG	A085583	001	
WHITEWORTH TOWN PLSN	200MG	A085444	001	

QUINORA

KEY PHARMS	200MG	A083576	001	
+ SCHERING	300MG	A085222	001	

TABLET, EXTENDED RELEASE;ORAL

QUINIDEX

WYETH PHARMS INC	300MG	N012796	002	
------------------	-------	---------	-----	--

QUINIDINE SULFATE

COSETTE	300MG	A040045	001	Jun 30, 1994
---------	-------	---------	-----	--------------

QUININE SULFATE

CAPSULE;ORAL

QUALAQUIN

+ SUN PHARM INDUSTRIES	324MG	N021799	001	Aug 12, 2005
------------------------	-------	---------	-----	--------------

QUININE SULFATE

AUROBINDO PHARMA USA	324MG	A202581	001	Dec 14, 2012
----------------------	-------	---------	-----	--------------

RABEPRAZOLE SODIUM

CAPSULE, DELAYED RELEASE;ORAL

ACIPHEX SPRINKLE

+ AYTU	5MG	N204736	001	Mar 26, 2013
+	10MG	N204736	002	Mar 26, 2013

TABLET, DELAYED RELEASE;ORAL

ACIPHEX

+ WAYLIS THERAP	10MG **	N020973	001	May 29, 2002
-----------------	---------	---------	-----	--------------

RABEPRAZOLE SODIUM

NORVIUM BIOSCIENCE	20MG	A076885	001	Nov 08, 2013
TEVA PHARMS USA	20MG	A076822	001	Nov 08, 2013

RAMIPRIL

CAPSULE;ORAL

RAMIPRIL

ACCORD HLTHCARE	1.25MG	A202392	001	Apr 15, 2014
	2.5MG	A202392	002	Apr 15, 2014
	5MG	A202392	003	Apr 15, 2014
	10MG	A202392	004	Apr 15, 2014
ACTAVIS ELIZABETH	1.25MG	A077513	001	Jun 18, 2008
	2.5MG	A077513	002	Jun 18, 2008
	5MG	A077513	003	Jun 18, 2008
	10MG	A077513	004	Jun 18, 2008
CHARTWELL RX	1.25MG	A077514	001	Jun 18, 2008
	2.5MG	A077514	002	Jun 18, 2008
	5MG	A077514	003	Jun 18, 2008
	10MG	A077514	004	Jun 18, 2008

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

RAMIPRIL

CAPSULE; ORAL

RAMIPRIL

CIPLA	1.25MG	A077004 001	Aug 07, 2008
	2.5MG	A077004 002	Aug 07, 2008
	5MG	A077004 003	Aug 07, 2008
	10MG	A077004 004	Aug 07, 2008
RANBAXY LABS LTD	5MG	A078849 001	Mar 06, 2009
	10MG	A078849 002	Mar 06, 2009
TEVA PHARMS	1.25MG	A077470 001	Jun 18, 2008
	2.5MG	A077470 002	Jun 18, 2008
	5MG	A077470 003	Jun 18, 2008
	10MG	A077470 004	Jun 18, 2008
WATSON LABS	5MG	A076549 003	Oct 24, 2005

TABLET; ORAL

ALTACE

+ KING PFIZER	1.25MG **	N022021 001	Feb 27, 2007
+	2.5MG **	N022021 002	Feb 27, 2007
+	5MG **	N022021 003	Feb 27, 2007
+	10MG **	N022021 004	Feb 27, 2007

RAMIPRIL

APOTEX	1.25MG	A091069 001	Dec 02, 2015
	2.5MG	A091069 002	Dec 02, 2015
	5MG	A091069 003	Dec 02, 2015
	10MG	A091069 004	Dec 02, 2015
NATCO PHARMA	1.25MG	A090650 001	Jun 30, 2011
	2.5MG	A090650 002	Jun 30, 2011
	5MG	A090650 003	Jun 30, 2011
	10MG	A090650 004	Jun 30, 2011
ZYDUS PHARMS USA INC	1.25MG	A090697 001	Sep 24, 2009
	2.5MG	A090697 002	Sep 24, 2009
	5MG	A090697 003	Sep 24, 2009
	10MG	A090697 004	Sep 24, 2009

RANITIDINE BISMUTH CITRATE

TABLET; ORAL

TRITEC

GLAXOSMITHKLINE	400MG	N020559 001	Aug 08, 1996
-----------------	-------	-------------	--------------

RANITIDINE HYDROCHLORIDE

CAPSULE; ORAL

RANITIDINE HYDROCHLORIDE

AJANTA PHARMA LTD	EQ 150MG BASE	A209859 001	Sep 27, 2018
	EQ 300MG BASE	A209859 002	Sep 27, 2018
APPCO	EQ 150MG BASE	A211893 001	Apr 05, 2019
	EQ 300MG BASE	A211893 002	Apr 05, 2019
AUROBINDO PHARMA	EQ 150MG BASE	A211058 001	Jul 16, 2018
	EQ 300MG BASE	A211058 002	Jul 16, 2018
NORVIUM BIOSCIENCE	EQ 150MG BASE	A075564 001	Oct 27, 2000
	EQ 300MG BASE	A075564 002	Oct 27, 2000
NOVITIUM PHARMA	EQ 150MG BASE	A210681 001	Nov 23, 2018
	EQ 300MG BASE	A210681 002	Nov 23, 2018
TEVA	EQ 150MG BASE	A075557 001	Oct 31, 2003
	EQ 300MG BASE	A075557 002	Oct 31, 2003

ZANTAC 150

+ GLAXOSMITHKLINE	EQ 150MG BASE **	N020095 001	Mar 08, 1994
-------------------	------------------	-------------	--------------

ZANTAC 300

+ GLAXOSMITHKLINE	EQ 300MG BASE **	N020095 002	Mar 08, 1994
-------------------	------------------	-------------	--------------

GRANULE, EFFERVESCENT; ORAL

ZANTAC 150

GLAXO GRP LTD	EQ 150MG BASE/PACKET	N020251 002	Mar 31, 1994
---------------	----------------------	-------------	--------------

INJECTABLE; INJECTION

RANITIDINE HYDROCHLORIDE

BEDFORD	EQ 25MG BASE/ML	A074764 001	Nov 19, 2004
HIKMA	EQ 25MG BASE/ML	A074777 001	Mar 02, 2005
	EQ 25MG BASE/ML	A077458 001	Feb 16, 2006
MYLAN LABS LTD	EQ 25MG BASE/ML	A079076 001	Jun 09, 2016
ZYDUS PHARMS USA INC	EQ 25MG BASE/ML	A091534 001	Feb 22, 2013

ZANTAC

+ PAI HOLDINGS PHARM	EQ 25MG BASE/ML	N019090 001	Oct 19, 1984
----------------------	-----------------	-------------	--------------

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

RANITIDINE HYDROCHLORIDE

INJECTABLE; INJECTION

ZANTAC IN PLASTIC CONTAINER

PAI HOLDINGS PHARM	EQ 1MG BASE/ML	N019593 002	Sep 27, 1991
	EQ 50MG BASE/100ML	N019593 001	Dec 17, 1986

SYRUP; ORAL

RANITIDINE HYDROCHLORIDE

ACTAVIS MID ATLANTIC	EQ 15MG BASE/ML	A076124 001	Feb 21, 2007
AMNEAL PHARMS	EQ 15MG BASE/ML	A078312 001	Sep 02, 2008
APOTEX INC	EQ 15MG BASE/ML	A077602 001	Sep 17, 2007
AUROBINDO PHARMA	EQ 15MG BASE/ML	A090623 001	Jul 28, 2010
EPIC PHARMA LLC	EQ 15MG BASE/ML	A091078 001	Mar 22, 2011
LANNETT CO INC	EQ 15MG BASE/ML	A078890 001	Jul 01, 2010
	EQ 15MG BASE/ML	A091288 001	Dec 09, 2010
NOSTRUM LABS INC	EQ 15MG BASE/ML	A078684 001	Aug 27, 2009
	EQ 15MG BASE/ML	A091091 001	Sep 20, 2011
PHARM ASSOC	EQ 15MG BASE/ML	A077405 001	Sep 21, 2007
RANBAXY	EQ 15MG BASE/ML	A078448 001	Dec 13, 2007
TARO	EQ 15MG BASE/ML	A077476 001	Jan 13, 2011
TOLMAR	EQ 15MG BASE/ML	A090054 001	Nov 15, 2010
TORRENT	EQ 15MG BASE/ML	A090102 001	May 26, 2009
WOCKHARDT	EQ 15MG BASE/ML	A079211 001	May 26, 2009
	EQ 15MG BASE/ML	A079212 001	Feb 23, 2009

ZANTAC

+ GLAXO GRP LTD

EQ 15MG BASE/ML N019675 001 Dec 30, 1988

TABLET; ORAL

RANITIDINE HYDROCHLORIDE

AMNEAL PHARMS NY	EQ 150MG BASE	A077824 001	Oct 13, 2006
	EQ 300MG BASE	A077824 002	Oct 13, 2006
ANI PHARMS	EQ 75MG BASE	A075212 001	Jan 14, 2000
	EQ 75MG BASE	A075296 001	Jan 14, 2000
	EQ 150MG BASE	A074488 001	Jul 31, 1997
	EQ 150MG BASE	A077426 001	Dec 19, 2005
	EQ 300MG BASE	A074488 002	Jul 31, 1997
	EQ 300MG BASE	A077426 002	Dec 19, 2005
APOTEX	EQ 75MG BASE	A075167 001	May 04, 2000
APOTEX INC	EQ 150MG BASE	A200172 001	May 31, 2012
AUROBINDO PHARMA	EQ 75MG BASE	A207579 001	Nov 13, 2017
	EQ 150MG BASE	A207578 001	Nov 13, 2017
BOEHRINGER INGELHEIM	EQ 150MG BASE	A074662 001	Aug 29, 1997
	EQ 300MG BASE	A074662 002	Aug 29, 1997
CONTRACT PHARMACAL	EQ 75MG BASE	A075094 001	Jun 21, 1999
ENDO OPERATIONS	EQ 150MG BASE	A075180 001	Jan 28, 1999
	EQ 300MG BASE	A075180 002	Jan 28, 1999
GRANULES	EQ 150MG BASE	A210243 001	Aug 20, 2018
	EQ 150MG BASE	A210243 002	Aug 20, 2018
HERITAGE PHARMA AVET	EQ 150MG BASE	A075165 001	Sep 30, 1998
	EQ 300MG BASE	A075165 002	Sep 30, 1998
MYLAN	EQ 150MG BASE	A074023 001	Aug 22, 1997
	EQ 300MG BASE	A074023 002	Aug 22, 1997
NORVIUM BIOSCIENCE	EQ 75MG BASE	A075497 001	Jan 14, 2000
	EQ 150MG BASE	A074552 001	Jul 30, 1998
	EQ 300MG BASE	A074552 002	Jul 30, 1998
NOSTRUM LABS INC	EQ 150MG BASE	A203694 001	Nov 30, 2017
	EQ 300MG BASE	A203694 002	Nov 30, 2017
PERRIGO	EQ 75MG BASE	A076195 001	Aug 30, 2002
PERRIGO R AND D	EQ 150MG BASE	A091429 001	May 11, 2011
	EQ 150MG BASE	A091429 002	May 11, 2011
RANBAXY	EQ 75MG BASE	A075254 001	Jan 14, 2000
	EQ 150MG BASE	A075000 001	Jan 30, 1998
	EQ 300MG BASE	A075000 002	Jan 30, 1998
SANDOZ	EQ 75MG BASE	A075519 001	Sep 26, 2002
STRIDES PHARMA	EQ 75MG BASE	A201745 001	Feb 29, 2012
	EQ 75MG BASE	A209160 001	Mar 05, 2018
	EQ 150MG BASE	A200536 001	Jun 28, 2011
	EQ 150MG BASE	A205512 001	Aug 22, 2016
	EQ 150MG BASE	A209161 001	Feb 22, 2018
	EQ 150MG BASE	A210010 001	Aug 01, 2018
	EQ 300MG BASE	A205512 002	Aug 22, 2016

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

RANITIDINE HYDROCHLORIDE

TABLET;ORAL

RANITIDINE HYDROCHLORIDE

	EQ 300MG BASE	A210010 002	Aug 01, 2018
SUN PHARM INDS LTD	EQ 75MG BASE	A075132 001	Jan 14, 2000
	EQ 150MG BASE	A075439 001	Apr 19, 2000
	EQ 300MG BASE	A075439 002	Apr 19, 2000
THINQ PHARM-CRO PVT	EQ 75MG BASE	A210250 001	Aug 30, 2019
	EQ 150MG BASE	A210228 001	Aug 30, 2019
WATSON LABS	EQ 150MG BASE	A074864 001	Oct 20, 1997
	EQ 300MG BASE	A074864 002	Oct 20, 1997
WOCKHARDT	EQ 75MG BASE	A076760 001	Feb 24, 2006
	EQ 75MG BASE	A078884 001	Jul 31, 2008
	EQ 150MG BASE	A078653 001	Nov 26, 2007
	EQ 150MG BASE	A078701 001	Nov 12, 2009
	EQ 300MG BASE	A078701 002	Dec 11, 2009
WOCKHARDT LTD	EQ 150MG BASE	A075208 001	Dec 17, 1998
	EQ 300MG BASE	A075208 002	Dec 17, 1998
ZANTAC 150			
+ CHATTEM SANOFI	EQ 150MG BASE	N021698 001	Aug 31, 2004
+	EQ 150MG BASE	N021698 002	Mar 13, 2007
+ GLAXO GRP LTD	EQ 150MG BASE **	N018703 001	Jun 09, 1983
ZANTAC 300			
+ GLAXO GRP LTD	EQ 300MG BASE **	N018703 002	Dec 09, 1985
ZANTAC 75			
+ CHATTEM SANOFI	EQ 75MG BASE	N020520 001	Dec 19, 1995
TABLET, EFFERVESCENT;ORAL			
ZANTAC 150			
GLAXO GRP LTD	EQ 150MG BASE	N020251 001	Mar 31, 1994
ZANTAC 25			
GLAXO GRP LTD	EQ 25MG BASE	N020251 003	Apr 01, 2004
ZANTAC 75			
+ CHATTEM SANOFI	EQ 75MG BASE **	N020745 001	Feb 26, 1998

RANOLAZINE

GRANULES, EXTENDED RELEASE;ORAL

ASPRUZYO SPRINKLE

+ SPIL	500MG	N216018 001	Feb 28, 2022
+	1GM	N216018 002	Feb 28, 2022

TABLET, EXTENDED RELEASE;ORAL

RANEXA

+ MENARINI INTL	500MG **	N021526 002	Jan 27, 2006
+	1GM **	N021526 001	Feb 12, 2007

RANOLAZINE

ACCORD HLTHCARE	500MG	A212930 001	May 18, 2021
	1GM	A212930 002	May 18, 2021
AMNEAL	500MG	A207690 001	Mar 11, 2021
	1GM	A207690 002	Mar 11, 2021
ANI PHARMS	500MG	A210482 001	Oct 29, 2019
	1GM	A210482 002	Oct 29, 2019
CIPLA	500MG	A211291 001	May 28, 2019
	1GM	A211291 002	May 28, 2019
PIRAMAL	500MG	A213085 001	Jul 25, 2023
	1GM	A213085 002	Jul 25, 2023

RAPACURONIUM BROMIDE

INJECTABLE;INJECTION

RAPLON

ORGANON USA INC	100MG/VIAL	N020984 001	Aug 18, 1999
	200MG/VIAL	N020984 002	Aug 18, 1999

RASAGILINE MESYLATE

TABLET;ORAL

RASAGILINE MESYLATE

APOTEX INC	EQ 0.5MG BASE	A201950 001	Sep 12, 2013
	EQ 1MG BASE	A201950 002	Sep 12, 2013
RYAN LABS	EQ 0.5MG BASE	A201942 001	Nov 18, 2021
	EQ 1MG BASE	A201942 002	Nov 18, 2021
WATSON LABS INC	EQ 0.5MG BASE	A201823 001	Jul 01, 2013
	EQ 1MG BASE	A201823 002	Jul 01, 2013

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

RAUWOLFIA SERPENTINA ROOT

TABLET;ORAL

HIWOLFIA

BOWMAN PHARMS	50MG	N009276 003
	50MG	N009276 005
	100MG	N009276 004

HYSERPIN

PHYS PRODS VA	50MG	N010581 001
---------------	------	-------------

KOGLUCOID

PANRAY	50MG	N009278 001
	100MG	N009278 002

RAUDIXIN

APOTHECON	50MG	N008842 001
	100MG	N008842 002

RAUSERPIN

FERNDAL LABS	50MG	N009926 002
	100MG	N009926 004

RAUVAL

PAL PAK	50MG	N009108 002
	100MG	N009108 004

RAUWOLFIA SERPENTINA

BUNDY	50MG	N009477 001
	100MG	N009477 002

HALSEY	50MG	A080498 001
	100MG	A080498 002

IMPAX LABS	50MG	N009273 001
	100MG	N009273 002

IVAX SUB TEVA PHARMS	50MG	N011521 001
	100MG	N011521 002

PUREPAC PHARM	50MG	A080842 001
	100MG	A080842 002

PVT FORM	50MG	A080583 001
	100MG	A080583 002

SOLVAY	50MG	A080500 001
	100MG	A080500 002

TABLICAPS	50MG	A083867 001
	100MG	A083444 001

VALEANT PHARM INTL	50MG	N009668 001
	100MG	N009668 002

WATSON LABS	50MG	A080907 001
	100MG	A080914 001

WOLFINA

FOREST PHARMS	50MG	N009255 008
	100MG	N009255 006

REPAGLINIDE

TABLET;ORAL

PRANDIN

+	GEMINI LABS LLC	0.5MG **	N020741 001	Dec 22, 1997
+		1MG **	N020741 002	Dec 22, 1997
+		2MG **	N020741 003	Dec 22, 1997

REPAGLINIDE

ACTAVIS TOTOWA	0.5MG	A090008 001	Jan 22, 2014
	1MG	A090008 002	Jan 22, 2014
	2MG	A090008 003	Jan 22, 2014

KENTON	0.5MG	A091517 001	Apr 24, 2015
	1MG	A091517 002	Apr 24, 2015
	2MG	A091517 003	Apr 24, 2015

NORVIUM BIOSCIENCE	0.5MG	A090252 001	Aug 23, 2013
	1MG	A090252 002	Jan 22, 2014
	2MG	A090252 003	Jan 22, 2014

RESCINNAMINE

CAPSULE;ORAL

CINNASIL

PANRAY	0.5MG	A084736 001
--------	-------	-------------

TABLET;ORAL

MODERIL

PFIZER	0.25MG	N010686 003
	0.5MG	N010686 006

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

RESERPINE

ELIXIR; ORAL

SERPASIL

NOVARTIS 0.2MG/4ML N009115 005

INJECTABLE; INJECTION

SANDRIL

LILLY 2.5MG/ML N010012 001

SERPASIL

NOVARTIS 2.5MG/ML N009434 002

TABLET; ORAL

HISERPIA

BOWMAN PHARMS 0.1MG N009631 002

0.25MG N009631 004

RAU-SED

BRISTOL MYERS SQUIBB 0.1MG N009357 001

0.25MG N009357 004

0.5MG N009357 006

1MG N009357 008

RESERPINE

BARR 0.25MG A080721 002

BELL PHARMA 0.1MG A083058 001

0.25MG A083058 002

BUNDY 0.1MG N009663 001

0.25MG N009663 003

CYCLE 0.1MG N009859 001

0.25MG N009859 002

ELKINS SINN 0.1MG A083145 001

0.25MG A083145 002

EVERYLIFE 0.1MG N010441 001

0.25MG N010441 002

0.5MG N010441 003

1MG N010441 004

HALSEY 0.1MG A080457 002

0.25MG A080457 001

1MG A080457 003

HIKMA INTL PHARMS 0.1MG A080975 001

0.25MG A080975 002

1MG A080975 003

IMPAX LABS 0.1MG N009627 001

0.25MG N009627 002

IVAX SUB TEVA PHARMS 0.1MG N011185 001

0.25MG N011185 002

MARSHALL PHARMA 0.1MG A080492 001

0.25MG A080492 002

MK LABS 0.1MG A080525 002

0.25MG A080525 001

MYLAN 1MG A084974 001

PHARMAVITE 0.25MG A084663 001

PUREPAC PHARM 0.1MG A080753 002

0.25MG A080753 001

PVT FORM 0.1MG A086117 001

0.25MG A080582 001

0.25MG A085775 001

1MG A080582 002

REXALL 0.25MG A080637 001

+ SANDOZ 0.1MG N009838 001

+ 0.25MG N009838 002

SOLVAY 0.25MG A080446 001

TABLICAPS 0.25MG A085207 001

TEVA 0.1MG A089020 001 Mar 07, 1985

0.25MG A089019 001 Mar 07, 1985

VALEANT PHARM INTL 0.1MG N009667 001

0.25MG N009667 002

WATSON LABS 0.1MG A080679 001

0.25MG A080393 001

0.25MG A085401 001

1MG A080749 001

WHITEWORTH TOWN PLSN 0.1MG A080723 001

0.25MG A080723 002

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

RESERPINE

TABLET;ORAL

RESERPINE

	1MG	A080723 003
SANDRIL		
LILLY	0.1MG	N009376 004
	0.25MG	N009376 001
SERPALAN		
LANNETT	0.1MG	N010124 001
	0.25MG	N010124 002
SERPANRAY		
PANRAY	0.1MG	N009391 001
	0.25MG	N009391 002
	1MG	N009391 004
SERPASIL		
NOVARTIS	0.1MG	N009115 001
	0.25MG	N009115 003
	1MG	N009115 004
SERPATE		
VALE	0.1MG	N009453 001
	0.25MG	N009453 002
SERPIVITE		
VITARINE	0.25MG	N009645 002

RESERPINE; TRICHLORMETHIAZIDE

TABLET;ORAL

METATENSIN #2

SANOFI AVENTIS US	0.1MG;2MG	N012972 001
METATENSIN #4		
SANOFI AVENTIS US	0.1MG;4MG	N012972 002
NAQUIVAL		
SCHERING	0.1MG;4MG	N012265 003
TRICHLORMETHIAZIDE W/ RESERPINE		
WATSON LABS	0.1MG;4MG	A085248 001

RETAPAMULIN

OINTMENT;TOPICAL

ALTABAX

+ ALMIRALL

1%

N022055 001 Apr 12, 2007

RIBAVIRIN

CAPSULE;ORAL

REBETOL

MERCK SHARP DOHME	200MG**Indicated for use and comarketed with Interferon ALFA-2B, Recombinant (INTRON A), as Rebetron Combination Therapy**	N020903 001 Jun 03, 1998
+	200MG **	N020903 002 Jul 25, 2001
RIBASPHERE		
CHARTWELL RX	200MG	A076203 001 Apr 06, 2004
RIBAVIRIN		
CHARTWELL RX	200MG	A076192 001 Apr 06, 2004
TEVA	200MG	A076277 001 Oct 04, 2004
SOLUTION;ORAL		
REBETOL		
+	SCHERING 40MG/ML	N021546 001 Jul 29, 2003
TABLET;ORAL		
COPEGUS		
+	ROCHE 200MG **	N021511 001 Dec 03, 2002
+	400MG **	N021511 002 Jun 21, 2005
RIBAVIRIN		
BEXIMCO PHARMS USA	200MG	A202546 001 Aug 12, 2014
	400MG	A202546 002 Aug 12, 2014
	500MG	A202546 003 Aug 12, 2014
	600MG	A202546 004 Aug 12, 2014
CHARTWELL RX	200MG	A077456 001 Dec 05, 2005
	400MG	A077456 002 Dec 05, 2005
	600MG	A077456 003 Dec 05, 2005
HERITAGE PHARMA AVET	200MG	A077053 001 Dec 05, 2005
ZYDUS PHARMS USA	400MG	A077094 002 Mar 16, 2007
	500MG	A077094 004 Apr 18, 2008

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

RIBAVIRINTABLET; ORAL
RIBAVIRIN

600MG

A077094 003 Mar 16, 2007

RIFAMPINCAPSULE; ORAL
RIFADIN

SANOFI AVENTIS US

150MG

A062303 001

+

300MG

N050420 001

RIFAMPIN

HIKMA

150MG

A065028 001 Mar 14, 2001

300MG

A065028 002 Mar 14, 2001

INJECTABLE; INJECTION

RIFAMPIN

AVET LIFESCIENCES

600MG/VIAL

A204101 001 Aug 18, 2014

WATSON PHARMS TEVA

600MG/VIAL

A206736 001 Jan 19, 2016

RIFAMYCIN SODIUM

TABLET, DELAYED RELEASE; ORAL

AEMCOLO

+

COSMO TECHNOLOGIES

EQ 194MG BASE

N210910 001 Nov 16, 2018

RILPIVIRINE HYDROCHLORIDE

TABLET, FOR SUSPENSION; ORAL

EDURANT PED

+

JANSSEN PRODS

EQ 2.5MG BASE

N219016 001 Mar 15, 2024

RILUZOLE

FILM; ORAL

EXSERVAN

+

AQUESTIVE

50MG

N212640 001 Nov 22, 2019

TABLET; ORAL

RILUTEK

+

COVIS

50MG

N020599 001 Dec 12, 1995

RILUZOLE

APOTEX CORP

50MG

A091300 001 Jun 18, 2013

DAITO PHARMS CO LTD

50MG

A204430 001 Oct 16, 2018

SUN PHARM INDS LTD

50MG

A091417 001 Jun 18, 2013

RIMANTADINE HYDROCHLORIDE

SYRUP; ORAL

FLUMADINE

FOREST LABS

50MG/5ML

N019650 001 Sep 17, 1993

TABLET; ORAL

FLUMADINE

+

SUN PHARM INDS INC

100MG

N019649 001 Sep 17, 1993

RIMANTADINE HYDROCHLORIDE

CHARTWELL RX

100MG

A076375 001 Jan 14, 2003

IMPAX LABS INC

100MG

A075916 001 Nov 02, 2001

RIMEXOLONE

SUSPENSION/DROPS; OPHTHALMIC

VEXOL

HARROW EYE

1%

N020474 001 Dec 30, 1994

RISEDRONATE SODIUM

TABLET; ORAL

ACTONEL

+

APIL

5MG

N020835 002 Apr 14, 2000

+

30MG

N020835 001 Mar 27, 1998

+

75MG **

N020835 004 Apr 16, 2007

RISEDRONATE SODIUM

HANGZHOU BINJIANG

35MG

A207516 001 Feb 15, 2019

NORVIUM BIOSCIENCE

5MG

A200477 001 Nov 30, 2015

30MG

A200477 002 Nov 30, 2015

35MG

A200477 003 Nov 30, 2015

75MG

A200477 004 Jun 10, 2014

150MG

A200477 005 Jun 10, 2014

TABLET, DELAYED RELEASE; ORAL

RISEDRONATE SODIUM

IMPAX LABS INC

35MG

A205066 001 Jun 29, 2018

ZYDUS PHARMS

35MG

A203822 001 Sep 11, 2018

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

RISPERIDONE

FOR SUSPENSION, EXTENDED RELEASE; INTRAMUSCULAR

RISVAN

+	LABS FARMS ROVI SA	75MG	N214835	001	Mar 29, 2024
+		100MG	N214835	002	Mar 29, 2024

SOLUTION; ORAL

RISPERIDONE

ANI PHARMS	1MG/ML	A076440	001	Jan 30, 2009
HIKMA	1MG/ML	A076904	001	Jul 29, 2009
PHARM ASSOC	1MG/ML	A077719	001	Jul 29, 2009
PRECISION DOSE	1MG/ML	A076797	001	Jun 28, 2010
SCIEGEN PHARMS INC	1MG/ML	A078909	001	Jul 29, 2009
WOCKHARDT	1MG/ML	A078744	001	Oct 08, 2009

TABLET; ORAL

RISPERDAL

JANSSEN PHARMS	5MG	N020272	005	Dec 29, 1993
----------------	-----	---------	-----	--------------

RISPERIDONE

DASH PHARMS	0.25MG	A076288	001	Sep 15, 2008
	0.5MG	A076288	002	Sep 15, 2008
	1MG	A076288	003	Sep 15, 2008
	2MG	A076288	004	Sep 15, 2008
	3MG	A076288	005	Sep 15, 2008
	4MG	A076288	006	Sep 15, 2008
HERITAGE PHARMA AVET	0.25MG	A076228	001	Jun 30, 2008
	0.25MG	A077769	001	Oct 16, 2008
	0.5MG	A076228	002	Jun 30, 2008
	0.5MG	A077769	002	Oct 16, 2008
	1MG	A076228	003	Jun 30, 2008
	1MG	A077769	003	Oct 16, 2008
	2MG	A076228	004	Jun 30, 2008
	2MG	A077769	004	Oct 16, 2008
	3MG	A076228	005	Jun 30, 2008
	3MG	A077769	005	Oct 16, 2008
	4MG	A076228	006	Jun 30, 2008
	4MG	A077769	006	Oct 16, 2008
JUBILANT CADISTA	0.25MG	A078828	001	Mar 23, 2009
	0.5MG	A078828	002	Mar 23, 2009
	1MG	A078828	003	Mar 23, 2009
	2MG	A078828	004	Mar 23, 2009
	3MG	A078828	005	Mar 23, 2009
	4MG	A078828	006	Mar 23, 2009
RATIOPHARM	0.25MG	A077784	001	Jun 08, 2010
	0.5MG	A077784	002	Jun 08, 2010
	1MG	A077784	003	Jun 08, 2010
	2MG	A077784	004	Jun 08, 2010
	3MG	A077784	005	Jun 08, 2010
	4MG	A077784	006	Jun 08, 2010
SUN PHARM INDS INC	0.25MG	A078036	001	Mar 10, 2014
	0.5MG	A078036	002	Mar 10, 2014
	1MG	A078036	003	Mar 10, 2014
	2MG	A078036	004	Mar 10, 2014
	3MG	A078036	005	Mar 10, 2014
	4MG	A078036	006	Mar 10, 2014
SYNTHON PHARMS	0.25MG	A078187	001	Oct 22, 2009
	0.5MG	A078187	002	Oct 22, 2009
	1MG	A078187	003	Oct 22, 2009
	2MG	A078187	004	Oct 22, 2009
	3MG	A078187	005	Oct 22, 2009
	4MG	A078187	006	Oct 22, 2009
WATSON LABS	0.25MG	A077860	001	Dec 05, 2008
	0.5MG	A077860	002	Dec 05, 2008
	1MG	A077860	003	Dec 05, 2008
	2MG	A077860	004	Dec 05, 2008
	3MG	A077860	005	Dec 05, 2008
	4MG	A077860	006	Dec 05, 2008
WEST WARD PHARMS	0.25MG	A078740	001	May 29, 2009
	0.5MG	A078740	002	May 29, 2009
	1MG	A078740	003	May 29, 2009
	2MG	A078740	004	May 29, 2009

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

RISPERIDONETABLET; ORAL
RISPERIDONE

3MG	A078740	005	May 29, 2009
4MG	A078740	006	May 29, 2009

TABLET, ORALLY DISINTEGRATING; ORAL

RISPERDAL

+	JANSSEN PHARMS	0.5MG **	N021444	001	Apr 02, 2003
+		1MG **	N021444	002	Apr 02, 2003
+		2MG **	N021444	003	Apr 02, 2003
+		3MG **	N021444	004	Dec 23, 2004
+		4MG **	N021444	005	Dec 23, 2004

RISPERIDONE

ACTAVIS LABS FL INC	0.5MG	A076996	001	Apr 19, 2011
	1MG	A076996	002	Apr 19, 2011
	2MG	A076996	003	Apr 19, 2011
	3MG	A076996	004	Apr 19, 2011
	4MG	A076996	005	Apr 19, 2011
CHARTWELL RX	0.5MG	A076908	001	Mar 12, 2012
	1MG	A076908	002	Mar 12, 2012
	2MG	A076908	003	Mar 12, 2012
DASH PHARMS	0.25MG	A091537	006	Feb 12, 2013
	0.5MG	A091537	001	Mar 30, 2011
	1MG	A091537	002	Mar 30, 2011
	2MG	A091537	003	Mar 30, 2011
	3MG	A091537	004	Mar 30, 2011
	4MG	A091537	005	Mar 30, 2011

RITODRINE HYDROCHLORIDE

INJECTABLE; INJECTION

RITODRINE HYDROCHLORIDE

ABRAXIS PHARM	10MG/ML	A071188	001	Jul 23, 1987
	15MG/ML	A071189	001	Jul 23, 1987
HOSPIRA	10MG/ML	A071618	001	Feb 28, 1991
	15MG/ML	A071619	001	Feb 28, 1991
RITODRINE HYDROCHLORIDE IN DEXTROSE 5% IN PLASTIC CONTAINER				
HOSPIRA	30MG/100ML	A071438	001	Jan 22, 1991

YUTOPAR

ASTRAZENECA	10MG/ML	N018580	001	
	15MG/ML	N018580	002	Sep 27, 1984

TABLET; ORAL

YUTOPAR

ASTRAZENECA	10MG	N018555	001	
-------------	------	---------	-----	--

RITONAVIR

CAPSULE; ORAL

NORVIR

ABBOTT	100MG	N020680	001	Mar 01, 1996	
+	ABBVIE	100MG **	N020945	001	Jun 29, 1999

RITONAVIR

HIKMA	100MG	A205801	001	Dec 03, 2020
-------	-------	---------	-----	--------------

SOLUTION; ORAL

NORVIR

+	ABBVIE	80MG/ML	N020659	001	Mar 01, 1996
---	--------	---------	---------	-----	--------------

RIVASTIGMINE TARTRATE

CAPSULE; ORAL

EXELON

+	NOVARTIS	EQ 1.5MG BASE **	N020823	003	Apr 21, 2000
+		EQ 3MG BASE **	N020823	004	Apr 21, 2000
+		EQ 4.5MG BASE **	N020823	005	Apr 21, 2000
+		EQ 6MG BASE **	N020823	006	Apr 21, 2000

RIVASTIGMINE TARTRATE

APOTEX INC	EQ 1.5MG BASE	A091072	001	May 16, 2013
	EQ 3MG BASE	A091072	002	May 16, 2013
	EQ 4.5MG BASE	A091072	003	May 16, 2013
	EQ 6MG BASE	A091072	004	May 16, 2013
SUN PHARM	EQ 1.5MG BASE	A077131	001	Oct 22, 2007
	EQ 3MG BASE	A077131	002	Oct 22, 2007
	EQ 4.5MG BASE	A077131	003	Oct 22, 2007
	EQ 6MG BASE	A077131	004	Oct 22, 2007

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

RIVASTIGMINE TARTRATE

SOLUTION;ORAL

EXELON

+ NOVARTIS

EQ 2MG BASE/ML

N021025 001 Apr 21, 2000

RIZATRIPTAN BENZOATE

FILM;ORAL

RIZAFILM

+ GENSCO

EQ 10MG BASE

N205394 001 Apr 14, 2023

TABLET;ORAL

MAXALT

+ ORGANON LLC

EQ 5MG BASE **

N020864 001 Jun 29, 1998

RIZATRIPTAN BENZOATE

APOTEX INC

EQ 5MG BASE

A202244 001 Dec 31, 2012

EQ 10MG BASE

A202244 002 Dec 31, 2012

AVET LIFESCIENCES

EQ 5MG BASE

A204090 001 Nov 26, 2013

EQ 10MG BASE

A204090 002 Nov 26, 2013

GLENMARK PHARMS LTD

EQ 5MG BASE

A201967 001 Dec 31, 2012

EQ 10MG BASE

A201967 002 Dec 31, 2012

INVAGEN PHARMS

EQ 5MG BASE

A204339 001 Jul 01, 2013

EQ 10MG BASE

A204339 002 Jul 01, 2013

JUBILANT GENERICS

EQ 5MG BASE

A203252 001 Dec 31, 2014

EQ 10MG BASE

A203252 002 Dec 31, 2014

NATCO PHARMA LTD

EQ 5MG BASE

A200482 001 Dec 31, 2012

EQ 10MG BASE

A200482 002 Dec 31, 2012

SANDOZ

EQ 5MG BASE

A079230 001 Dec 31, 2012

EQ 10MG BASE

A079230 002 Dec 31, 2012

UNICHEM

EQ 5MG BASE

A207836 001 Mar 07, 2017

EQ 10MG BASE

A207836 002 Mar 07, 2017

TABLET, ORALLY DISINTEGRATING;ORAL

MAXALT-MLT

+ ORGANON

EQ 5MG BASE **

N020865 001 Jun 29, 1998

RIZATRIPTAN BENZOATE

APOTEX INC

EQ 5MG BASE

A202477 001 Jul 01, 2013

EQ 10MG BASE

A202477 002 Jul 01, 2013

CHARTWELL RX

EQ 5MG BASE

A078739 001 Jul 01, 2013

EQ 10MG BASE

A078739 002 Jul 01, 2013

GLENMARK PHARMS LTD

EQ 5MG BASE

A201914 001 Jul 01, 2013

EQ 10MG BASE

A201914 002 Jul 01, 2013

JUBILANT GENERICS

EQ 5MG BASE

A203334 001 Oct 16, 2015

EQ 10MG BASE

A203334 002 Oct 16, 2015

NORVIUM BIOSCIENCE

EQ 5MG BASE

A078173 001 Dec 31, 2012

EQ 10MG BASE

A078173 002 Dec 31, 2012

UNICHEM

EQ 5MG BASE

A207835 001 Mar 07, 2017

EQ 10MG BASE

A207835 002 Mar 07, 2017

ROCURONIUM BROMIDE

INJECTABLE;INJECTION

ROCURONIUM BROMIDE

HIKMA

50MG/5ML (10MG/ML)

A217034 001 May 24, 2023

NORVIUM BIOSCIENCE

50MG/5ML (10MG/ML)

A204918 001 Jul 12, 2024

100MG/10ML (10MG/ML)

A204918 002 Jul 12, 2024

TEVA PHARMS

50MG/5ML (10MG/ML)

A078717 001 Nov 26, 2008

100MG/10ML (10MG/ML)

A078717 002 Nov 26, 2008

ZEMURON

+ ORGANON USA INC

50MG/5ML (10MG/ML) **

N020214 001 Mar 17, 1994

+

10MG/ML (10MG/ML) **

N020214 002 Mar 17, 1994

+

100MG/10ML (10MG/ML) **

N020214 003 Mar 17, 1994

ROFECOXIB

SUSPENSION;ORAL

VIOXX

MERCK

12.5MG/5ML

N021052 001 May 20, 1999

25MG/5ML

N021052 002 May 20, 1999

TABLET;ORAL

VIOXX

MERCK

12.5MG

N021042 001 May 20, 1999

25MG

N021042 002 May 20, 1999

50MG

N021042 003 Feb 25, 2000

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ROFLUMILAST

TABLET; ORAL

ROFLUMILAST

BRECKENRIDGE	250MCG	A208236 002	Oct 19, 2023
	500MCG	A208236 001	Oct 03, 2018
TORRENT	500MCG	A208272 001	Aug 06, 2018

ROLAPITANT HYDROCHLORIDE

EMULSION; INTRAVENOUS

VARUBI

+ TERSERA	EQ 166.5MG BASE/92.5ML (EQ 1.8MG BASE/ML)	N208399 001	Oct 25, 2017
-----------	---	-------------	--------------

ROMIDEPSIN

SOLUTION; INTRAVENOUS

ROMIDEPSIN

+ TEVA PHARMS USA INC	10MG/2ML (5MG/ML) **	N208574 001	Mar 13, 2020
+	27.5MG/5.5ML (5MG/ML) **	N208574 002	Mar 13, 2020

ROPINIROLE HYDROCHLORIDE

TABLET; ORAL

REQUIP

+ GLAXOSMITHKLINE LLC	EQ 0.25MG BASE **	N020658 001	Sep 19, 1997
+	EQ 0.5MG BASE **	N020658 002	Sep 19, 1997
+	EQ 1MG BASE **	N020658 003	Sep 19, 1997
+	EQ 2MG BASE **	N020658 004	Sep 19, 1997
+	EQ 3MG BASE **	N020658 006	Jan 27, 1999
+	EQ 4MG BASE **	N020658 007	Jan 27, 1999
+	EQ 5MG BASE **	N020658 005	Sep 19, 1997

ROPINIROLE HYDROCHLORIDE

COSETTE

	EQ 0.25MG BASE	A077460 001	May 05, 2008
	EQ 0.5MG BASE	A077460 002	May 05, 2008
	EQ 1MG BASE	A077460 003	May 05, 2008
	EQ 2MG BASE	A077460 004	May 05, 2008
	EQ 3MG BASE	A077460 005	May 05, 2008
	EQ 4MG BASE	A077460 006	May 05, 2008
	EQ 5MG BASE	A077460 007	May 19, 2008

EPIC PHARMA LLC

	EQ 0.25MG BASE	A078230 001	May 20, 2008
	EQ 0.5MG BASE	A078230 002	May 20, 2008
	EQ 1MG BASE	A078230 003	May 20, 2008
	EQ 2MG BASE	A078230 004	May 20, 2008
	EQ 3MG BASE	A078230 005	May 20, 2008
	EQ 4MG BASE	A078230 006	May 20, 2008
	EQ 5MG BASE	A078230 007	May 20, 2008

HIKMA

	EQ 0.25MG BASE	A077852 001	May 05, 2008
	EQ 0.5MG BASE	A077852 002	May 05, 2008
	EQ 1MG BASE	A077852 003	May 05, 2008
	EQ 2MG BASE	A077852 004	May 05, 2008
	EQ 3MG BASE	A077852 005	May 05, 2008
	EQ 4MG BASE	A077852 006	May 05, 2008
	EQ 5MG BASE	A077852 007	May 19, 2008

NORVIUM BIOSCIENCE

	EQ 0.25MG BASE	A078881 001	May 05, 2008
	EQ 0.5MG BASE	A078881 002	May 05, 2008
	EQ 1MG BASE	A078881 003	May 05, 2008
	EQ 2MG BASE	A078881 004	May 05, 2008
	EQ 3MG BASE	A078881 005	May 05, 2008
	EQ 4MG BASE	A078881 006	May 05, 2008
	EQ 5MG BASE	A078881 007	May 19, 2008

TABLET, EXTENDED RELEASE; ORAL

REQUIP XL

+ GLAXOSMITHKLINE LLC	EQ 2MG BASE **	N022008 001	Jun 13, 2008
+	EQ 3MG BASE **	N022008 002	Jun 13, 2008
+	EQ 4MG BASE **	N022008 003	Jun 13, 2008
+	EQ 6MG BASE **	N022008 006	Apr 10, 2009
+	EQ 8MG BASE **	N022008 004	Jun 13, 2008
+	EQ 12MG BASE **	N022008 005	Oct 31, 2008

ROPINIROLE HYDROCHLORIDE

MYLAN PHARMS INC

	EQ 2MG BASE	A200462 001	Oct 15, 2012
	EQ 3MG BASE	A200462 002	Oct 15, 2012
	EQ 4MG BASE	A200462 003	Oct 15, 2012
	EQ 6MG BASE	A200462 004	Oct 15, 2012

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ROPINIROLE HYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL

ROPINIROLE HYDROCHLORIDE

	EQ 8MG BASE	A200462 005	Oct 15, 2012
	EQ 12MG BASE	A200462 006	Oct 15, 2012
WATSON LABS INC	EQ 2MG BASE	A200431 001	Jun 06, 2012
	EQ 4MG BASE	A200431 002	Jun 06, 2012
	EQ 6MG BASE	A200431 003	Jun 06, 2012
	EQ 8MG BASE	A200431 004	Jun 06, 2012
	EQ 12MG BASE	A200431 005	Jun 06, 2012

ROPIVACAINE HYDROCHLORIDE

SOLUTION;INJECTION

NAROPIN

+	FRESENIUS KABI USA	50MG/10ML (5MG/ML) **	N020533 013	May 01, 1998
+		75MG/10ML (7.5MG/ML) **	N020533 012	Sep 24, 1996

ROPIVACAINE HYDROCHLORIDE

	HOSPIRA	20MG/10ML (2MG/ML)	A090194 001	Sep 23, 2014
		40MG/20ML (2MG/ML)	A090194 005	Sep 23, 2014
		100MG/10ML (10MG/ML)	A090194 004	Sep 23, 2014
		150MG/30ML (5MG/ML)	A090194 002	Sep 23, 2014
		150MG/20ML (7.5MG/ML)	A090194 003	Sep 23, 2014
		200MG/20ML (10MG/ML)	A090194 006	Sep 23, 2014
RISING		40MG/20ML (2MG/ML)	A090318 001	Sep 23, 2014
		150MG/30ML (5MG/ML)	A090318 002	Sep 23, 2014
		150MG/20ML (7.5MG/ML)	A090318 003	Sep 23, 2014
		200MG/20ML (10MG/ML)	A090318 004	Sep 23, 2014

ROSE BENGAL SODIUM I-131

INJECTABLE;INJECTION

ROBENGATOPE

	BRACCO	0.5mCi/VIAL	N016224 001
		1mCi/VIAL	N016224 002
		2mCi/VIAL	N016224 003

SODIUM ROSE BENGAL I 131

	SORIN	0.5mCi/ML	N017318 001
--	-------	-----------	-------------

ROSIGLITAZONE MALEATE

TABLET;ORAL

AVANDIA

+	WOODWARD	EQ 2MG BASE	N021071 002	May 25, 1999
+		EQ 4MG BASE	N021071 003	May 25, 1999
+		EQ 8MG BASE **	N021071 004	May 25, 1999

ROSIGLITAZONE MALEATE

	ANI PHARMS	EQ 2MG BASE	A076747 001	Jan 25, 2013
		EQ 4MG BASE	A076747 002	Jan 25, 2013
		EQ 8MG BASE	A076747 003	Jan 25, 2013

ROSUVASTATIN CALCIUM

CAPSULE;ORAL

EZALLOR SPRINKLE

+	SUN PHARM	EQ 5MG BASE	N208647 001	Dec 18, 2018
+		EQ 10MG BASE	N208647 002	Dec 18, 2018
+		EQ 20MG BASE	N208647 003	Dec 18, 2018
+		EQ 40MG BASE	N208647 004	Dec 18, 2018

TABLET;ORAL

ROSUVASTATIN CALCIUM

	AMNEAL PHARMS CO	EQ 5MG BASE	A208850 001	Oct 16, 2018
		EQ 10MG BASE	A208850 002	Oct 16, 2018
		EQ 20MG BASE	A208850 003	Oct 16, 2018
		EQ 40MG BASE	A208850 004	Oct 16, 2018
	APOTEX	EQ 5MG BASE	A079145 001	Jul 19, 2016
		EQ 10MG BASE	A079145 002	Jul 19, 2016
		EQ 20MG BASE	A079145 003	Jul 19, 2016
		EQ 40MG BASE	A079145 004	Jul 19, 2016
	INVENTIA	EQ 5MG BASE	A207653 001	Feb 05, 2021
		EQ 10MG BASE	A207653 002	Feb 05, 2021
		EQ 20MG BASE	A207653 003	Feb 05, 2021
		EQ 40MG BASE	A207653 004	Feb 05, 2021
	SANDOZ	EQ 5MG BASE	A079171 001	Jul 19, 2016
		EQ 10MG BASE	A079171 002	Jul 19, 2016

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ROSUVASTATIN CALCIUM

TABLET; ORAL

ROSUVASTATIN CALCIUM

	EQ 20MG BASE	A079171 003	Jul 19, 2016
	EQ 40MG BASE	A079171 004	Jul 19, 2016
STRIDES PHARMA	EQ 5MG BASE	A079161 001	Jul 19, 2016
	EQ 10MG BASE	A079161 002	Jul 19, 2016
	EQ 20MG BASE	A079161 003	Jul 19, 2016
	EQ 40MG BASE	A079161 004	Jul 19, 2016
SUN PHARM	EQ 5MG BASE	A079169 001	Jul 19, 2016
	EQ 10MG BASE	A079169 002	Jul 19, 2016
	EQ 20MG BASE	A079169 003	Jul 19, 2016
	EQ 40MG BASE	A079169 004	Jul 19, 2016
SUNSHINE	EQ 5MG BASE	A210667 001	Apr 01, 2020
	EQ 10MG BASE	A210667 002	Apr 01, 2020
	EQ 20MG BASE	A210667 003	Apr 01, 2020
	EQ 40MG BASE	A210667 004	Apr 01, 2020
TEVA PHARMS USA	EQ 5MG BASE	A079166 001	Jul 19, 2016
	EQ 10MG BASE	A079166 002	Jul 19, 2016
	EQ 20MG BASE	A079166 003	Jul 19, 2016
	EQ 40MG BASE	A079166 004	Jul 19, 2016
ZHEJIANG JINGXIN	EQ 5MG BASE	A206513 001	Mar 01, 2019
	EQ 10MG BASE	A206513 002	Mar 01, 2019
	EQ 20MG BASE	A206513 003	Mar 01, 2019
	EQ 40MG BASE	A206513 004	Mar 01, 2019

RUFINAMIDE

TABLET; ORAL

BANZEL

+ EISAI INC

100MG **

N021911 001 Nov 14, 2008

SAFFLOWER OIL

INJECTABLE; INJECTION

LIPOSYN 10%

ABBOTT

10% (10GM/100ML)

N018203 001

LIPOSYN 20%

ABBOTT

20% (20GM/100ML)

N018614 001

SAFFLOWER OIL; SOYBEAN OIL

INJECTABLE; INJECTION

LIPOSYN II 10%

HOSPIRA

5%;5% (5GM/100ML)

N018997 001 Aug 27, 1984

LIPOSYN II 20%

HOSPIRA

10%;10% (10GM/100ML)

N018991 001 Aug 27, 1984

SAFINAMIDE MESYLATE

TABLET; ORAL

SAFINAMIDE MESYLATE

AUROBINDO PHARMA

EQ 50MG BASE

A215902 001 Jun 14, 2023

EQ 100MG BASE

A215902 002 Jun 14, 2023

SALMETEROL XINAFOATE

AEROSOL, METERED; INHALATION

SEREVENT

GLAXOSMITHKLINE

EQ 0.021MG BASE/INH

N020236 001 Feb 04, 1994

SAMARIUM SM-153 LEXIDRONAM PENTASODIUM

INJECTABLE; INJECTION

QUADRAMET

+ LANTHEUS MEDICAL

50mCi/ML

N020570 001 Mar 28, 1997

SAQUINAVIR

CAPSULE; ORAL

FORTOVASE

+ HOFFMANN LA ROCHE

200MG **

N020828 001 Nov 07, 1997

SAQUINAVIR MESYLATE

CAPSULE; ORAL

INVIRASE

+ HOFFMANN LA ROCHE

EQ 200MG BASE

N020628 001 Dec 06, 1995

TABLET; ORAL

INVIRASE

+ HOFFMANN-LA ROCHE

EQ 500MG BASE

N021785 001 Dec 17, 2004

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

SARALASIN ACETATE

INJECTABLE; INJECTION

SARENIN

PROCTER AND GAMBLE EQ 0.6MG BASE/ML N018009 001

SAXAGLIPTIN HYDROCHLORIDE

TABLET; ORAL

ONGLYZA

+ ASTRAZENECA AB EQ 2.5MG BASE ** N022350 001 Jul 31, 2009

+ EQ 5MG BASE ** N022350 002 Jul 31, 2009

SAXAGLIPTIN

AMNEAL EQ 2.5MG BASE A205941 001 Jul 31, 2023

EQ 5MG BASE A205941 002 Jul 31, 2023

SECOBARBITAL SODIUM

CAPSULE; ORAL

SECOBARBITAL SODIUM

ANABOLIC 100MG A084422 001

BARR 100MG A084225 001

EVERYLIFE 100MG A085895 001

HALSEY 100MG A084676 001

IVAX PHARMS 100MG A085869 001

KV PHARM 100MG A085285 001

LANNETT 50MG A085909 001

100MG A085903 001

PARKE DAVIS 100MG A084762 001

PERRIGO 100MG A084561 001

PUREPAC PHARM 100MG A085867 001

VALEANT PHARM INTL 100MG A085477 001

VITARINE 100MG A085898 001

100MG A086273 001

WATSON LABS 100MG A085792 001

WEST WARD 100MG A084926 001

WHITWORTH TOWN PLSN 100MG A085798 001

WYETH AYERST 100MG A086390 001

SECONAL SODIUM

VALEANT PHARMS NORTH 50MG A086101 001 Oct 03, 1983

100MG A086101 002 Oct 03, 1983

INJECTABLE; INJECTION

SECOBARBITAL SODIUM

ELKINS SINN 100MG/VIAL A083281 001

WYETH AYERST 50MG/ML A083262 001

SECONAL SODIUM

LILLY 50MG/ML ** N007392 002

SUPPOSITORY; RECTAL

SECONAL SODIUM

LILLY 30MG A086530 001

60MG A086530 002

120MG A086530 003

200MG A086530 004

SECRETIN

INJECTABLE; INJECTION

SECRETIN-FERRING

FERRING 75CU/VIAL N018290 001

SECRETIN SYNTHETIC PORCINE

FOR SOLUTION; INTRAVENOUS

SECREFLO

CHIRHOCLIN 16MCG/VIAL N021136 001 Apr 04, 2002

SELEGILINE HYDROCHLORIDE

CAPSULE; ORAL

ELDEPRYL

+ NORVIUM BIOSCIENCE 5MG ** N020647 001 May 15, 1996

SELEGILINE HYDROCHLORIDE

LANNETT CO INC 5MG A075145 001 Sep 15, 2003

TABLET; ORAL

SELEGILINE HYDROCHLORIDE

CHARTWELL MOLECULES 5MG A074565 001 Aug 02, 1996

5MG A074641 001 Aug 02, 1996

COSETTE 5MG A074744 001 Jan 27, 1997

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

SELEGILINE HYDROCHLORIDE

TABLET; ORAL

SELEGILINE HYDROCHLORIDE

	5MG	A074756 001	Nov 25, 1998
ESJAY PHARMA	5MG	A074866 001	Nov 26, 1997
G AND W LABS INC	5MG	A074537 001	Aug 02, 1996
KENTON	5MG	A074912 001	Apr 30, 1998
+ NORVIUM BIOSCIENCE	5MG **	N019334 001	Jun 05, 1989

SELENIUM SULFIDE

LOTION/SHAMPOO; TOPICAL

EXSEL

ALLERGAN HERBERT	2.5%	A083892 001	
SELENIUM SULFIDE			
ACTAVIS MID ATLANTIC	2.5%	A084394 001	
COSETTE	2.5%	A086209 001	
IVAX PHARMS	2.5%	A085777 001	
PHARMOBEDIANT CNSLTG	2.5%	A088228 001	Sep 01, 1983
SELSUN			
+ CHATTEM	2.5% **	N007936 001	

SELENOMETHIONINE SE-75

INJECTABLE; INJECTION

SELENOMETHIONINE SE 75

GE HEALTHCARE	250uCi/ML	N017257 001	
MALLINCKRODT	100uCi/ML	N017098 001	
PHARMALUCENCE	500uCi/ML	N017322 001	
SETHOTOPE			
BRACCO	85-550uCi/ML	N017047 001	

SELEXIPAG

TABLET; ORAL

SELEXIPAG

ZYDUS LIFESCIENCES	0.2MG	A214302 001	Dec 21, 2022
	0.4MG	A214302 002	Dec 21, 2022
	0.6MG	A214302 003	Dec 21, 2022
	0.8MG	A214302 004	Dec 21, 2022
	1MG	A214302 005	Dec 21, 2022
	1.2MG	A214302 006	Dec 21, 2022
	1.4MG	A214302 007	Dec 21, 2022
	1.6MG	A214302 008	Dec 21, 2022

SERMORELIN ACETATE

INJECTABLE; INJECTION

GEREF

+ EMD SERONO	EQ 0.05MG BASE/AMP **	N019863 001	Dec 28, 1990
+ EMD SERONO INC	EQ 0.5MG BASE/VIAL **	N020443 001	Sep 26, 1997
+	EQ 1MG BASE/VIAL **	N020443 002	Sep 26, 1997

SERTRALINE HYDROCHLORIDE

CONCENTRATE; ORAL

SERTRALINE HYDROCHLORIDE

RANBAXY LABS LTD	EQ 20MG BASE/ML	A078053 001	Feb 05, 2007
------------------	-----------------	-------------	--------------

TABLET; ORAL

SERTRALINE HYDROCHLORIDE

ANDA REPOSITORY	EQ 25MG BASE	A077818 001	Feb 06, 2007
	EQ 50MG BASE	A077818 002	Feb 06, 2007
	EQ 100MG BASE	A077818 003	Feb 06, 2007
APPCO	EQ 25MG BASE	A077713 001	Feb 06, 2007
	EQ 50MG BASE	A077713 002	Feb 06, 2007
	EQ 100MG BASE	A077713 003	Feb 06, 2007
CHARTWELL MOLECULAR	EQ 25MG BASE	A077162 001	Feb 06, 2007
	EQ 50MG BASE	A077162 002	Feb 06, 2007
	EQ 100MG BASE	A077162 003	Feb 06, 2007
HERITAGE PHARMA AVET	EQ 25MG BASE	A077299 001	Feb 06, 2007
	EQ 25MG BASE	A077345 001	Feb 06, 2007
	EQ 25MG BASE	A077663 001	Feb 06, 2007
	EQ 50MG BASE	A077299 002	Feb 06, 2007
	EQ 50MG BASE	A077345 002	Feb 06, 2007
	EQ 50MG BASE	A077663 002	Feb 06, 2007
	EQ 100MG BASE	A077299 003	Feb 06, 2007
	EQ 100MG BASE	A077345 003	Feb 06, 2007

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

SERTRALINE HYDROCHLORIDE

TABLET;ORAL

SERTRALINE HYDROCHLORIDE

	EQ 100MG BASE	A077663 003	Feb 06, 2007
HIKMA PHARMS	EQ 25MG BASE	A077864 001	Aug 10, 2009
	EQ 50MG BASE	A077864 002	Aug 10, 2009
	EQ 100MG BASE	A077864 003	Aug 10, 2009
IVAX SUB TEVA PHARMS	EQ 25MG BASE	A075719 003	Jun 30, 2006
	EQ 50MG BASE	A075719 001	Jun 30, 2006
	EQ 100MG BASE	A075719 002	Jun 30, 2006
MYLAN PHARMS INC	EQ 25MG BASE	A076540 001	Mar 20, 2007
	EQ 50MG BASE	A076540 002	Mar 20, 2007
	EQ 100MG BASE	A076540 003	Mar 20, 2007
NORVIUM BIOSCIENCE	EQ 25MG BASE	A076671 001	Feb 06, 2007
	EQ 25MG BASE	A078626 001	Jan 31, 2008
	EQ 50MG BASE	A076671 002	Feb 06, 2007
	EQ 50MG BASE	A078626 002	Jan 31, 2008
	EQ 100MG BASE	A076671 003	Feb 06, 2007
	EQ 100MG BASE	A078626 003	Jan 31, 2008
SUN PHARM INDS (IN)	EQ 25MG BASE	A078108 001	Feb 06, 2007
	EQ 50MG BASE	A078108 002	Feb 06, 2007
	EQ 100MG BASE	A078108 003	Feb 06, 2007
SUN PHARM INDS LTD	EQ 25MG BASE	A077977 001	Feb 06, 2007
	EQ 50MG BASE	A077977 002	Feb 06, 2007
	EQ 100MG BASE	A077977 003	Feb 06, 2007
	EQ 150MG BASE	A077977 004	Feb 06, 2007
	EQ 200MG BASE	A077977 005	Feb 06, 2007
TORRENT PHARMS	EQ 25MG BASE	A077765 001	Feb 06, 2007
	EQ 50MG BASE	A077765 002	Feb 06, 2007
	EQ 100MG BASE	A077765 003	Feb 06, 2007
ZYDUS	EQ 25MG BASE	A077106 001	Feb 06, 2007
	EQ 50MG BASE	A077106 002	Feb 06, 2007
	EQ 100MG BASE	A077106 003	Feb 06, 2007
ZOLOFT			
+ VIATRIS	EQ 150MG BASE **	N019839 003	Dec 30, 1991
+	EQ 200MG BASE **	N019839 004	Dec 30, 1991

SEVELAMER CARBONATE

FOR SUSPENSION;ORAL

SEVELAMER CARBONATE

LUPIN LTD

800MG/PACKET

A201513 001

Dec 23, 2021

2.4GM/PACKET

A201513 002

Dec 23, 2021

TABLET;ORAL

SEVELAMER CARBONATE

CHARTWELL RX

800MG

A204600 001

Jan 14, 2021

IMPAX LABS INC

800MG

A090975 001

Oct 23, 2017

SINOTHERAPEUTICS INC

800MG

A212970 001

Apr 09, 2020

STRIDES PHARMA

800MG

A201069 001

Aug 05, 2020

SEVELAMER HYDROCHLORIDE

CAPSULE;ORAL

RENAGEL

GENZYME

403MG

N020926 001

Oct 30, 1998

TABLET;ORAL

SEVELAMER HYDROCHLORIDE

RISING

400MG

A201068 001

Dec 14, 2020

800MG

A201068 002

Dec 14, 2020

SIBUTRAMINE HYDROCHLORIDE

CAPSULE;ORAL

MERIDIA

ABBOTT

5MG

N020632 001

Nov 22, 1997

10MG

N020632 002

Nov 22, 1997

15MG

N020632 003

Nov 22, 1997

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

SILDENAFIL CITRATE

FOR SUSPENSION;ORAL

SILDENAFIL CITRATE

GRANULES

EQ 10MG BASE/ML

A214556 001 Dec 01, 2023

TRIS PHARMA INC

EQ 10MG BASE/ML

A212312 001 Nov 17, 2021

SUSPENSION;ORAL

LIQREV

+ CMP DEV LLC

EQ 10MG BASE/ML

N214952 001 Apr 28, 2023

TABLET;ORAL

SILDENAFIL CITRATE

ACTAVIS GRP PTC

EQ 20MG BASE

A200149 001 Feb 25, 2013

APOTEX CORP

EQ 20MG BASE

A091379 001 Nov 06, 2012

MYLAN PHARMS INC

EQ 20MG BASE

A201150 001 Nov 09, 2012

PERRIGO R AND D

EQ 25MG BASE

A205791 001 Apr 23, 2020

EQ 50MG BASE

A205791 002 Apr 23, 2020

WATSON LABS INC

EQ 25MG BASE

A202506 001 Nov 25, 2020

EQ 50MG BASE

A202506 002 Nov 25, 2020

EQ 100MG BASE

A202506 003 Nov 25, 2020

SILODOSIN

CAPSULE;ORAL

SILODOSIN

ALEMBIC

4MG

A211731 001 Nov 22, 2019

8MG

A211731 002 Nov 22, 2019

ZYDUS PHARMS

4MG

A204816 001 Dec 08, 2022

8MG

A204816 002 Dec 08, 2022

SILVER SULFADIAZINE

CREAM;TOPICAL

SSD AF

DR REDDYS

1%

N018578 003 Jul 11, 1990

DRESSING;TOPICAL

SILDAFLO

FRANKLIN PHARMS

1%

N019608 001 Nov 30, 1989

SIMEPREVIR SODIUM

CAPSULE;ORAL

OLYSIO

+ JANSSEN PRODS

EQ 150MG BASE

N205123 001 Nov 22, 2013

SIMETHICONE-CELLULOSE

SUSPENSION;ORAL

SONORX

BRACCO

7.5MG/ML

N020773 001 Oct 29, 1998

SIMVASTATIN

TABLET;ORAL

SIMVASTATIN

CHARTWELL RX

5MG

A077766 001 Dec 20, 2006

10MG

A077766 002 Dec 20, 2006

20MG

A077766 003 Dec 20, 2006

40MG

A077766 004 Dec 20, 2006

80MG

A077766 005 Dec 20, 2006

IVAX SUB TEVA PHARMS

5MG

A076052 001 Jun 23, 2006

10MG

A076052 002 Jun 23, 2006

20MG

A076052 003 Jun 23, 2006

40MG

A076052 004 Jun 23, 2006

80MG

A076052 005 Dec 20, 2006

NORVIUM BIOSCIENCE

5MG

A090868 001 Jun 08, 2010

10MG

A090868 002 Jun 08, 2010

20MG

A090868 003 Jun 08, 2010

40MG

A090868 004 Jun 08, 2010

80MG

A090868 005 Jun 08, 2010

SUN PHARM INDS LTD

5MG

A076285 001 Dec 20, 2006

10MG

A076285 002 Dec 20, 2006

20MG

A076285 003 Dec 20, 2006

40MG

A076285 004 Dec 20, 2006

80MG

A076285 005 Jun 23, 2006

WATSON LABS TEVA

5MG

A076685 001 Dec 20, 2006

10MG

A076685 002 Dec 20, 2006

20MG

A076685 003 Dec 20, 2006

40MG

A076685 004 Dec 20, 2006

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

SIMVASTATINTABLET; ORAL
SIMVASTATIN

	80MG	A076685 005	Dec 20, 2006
ZOCOR			
+ ORGANON	80MG **	N019766 005	Jul 10, 1998
TABLET, ORALLY DISINTEGRATING; ORAL			
SIMVASTATIN			
+ SYNTHON PHARMS	10MG	N021961 001	Oct 09, 2007
+	20MG	N021961 002	Oct 09, 2007
+	40MG	N021961 003	Oct 09, 2007
+	80MG	N021961 004	Oct 09, 2007

SIMVASTATIN; SITAGLIPTIN PHOSPHATE

TABLET; ORAL

JUVISYNC			
+ MERCK SHARP DOHME	10MG;EQ 50MG BASE **	N202343 004	Sep 18, 2012
+	10MG;EQ 100MG BASE **	N202343 001	Oct 07, 2011
+	20MG;EQ 50MG BASE **	N202343 005	Sep 18, 2012
+	20MG;EQ 100MG BASE **	N202343 002	Oct 07, 2011
+	40MG;EQ 50MG BASE **	N202343 006	Sep 18, 2012
+	40MG;EQ 100MG BASE **	N202343 003	Oct 07, 2011

SIROLIMUS

SOLUTION; ORAL

SIROLIMUS			
TORRENT	1MG/ML	A215016 001	Dec 27, 2021

TABLET; ORAL

RAPAMUNE			
+ PF PRISM CV	5MG **	N021110 003	Feb 23, 2004

SODIUM BENZOATE; SODIUM PHENYLACETATE

SOLUTION; INTRAVENOUS

AMMONUL			
+ BAUSCH	10%;10% (5GM/50ML;5GM/50ML)	N020645 001	Feb 17, 2005

SOLUTION; ORAL

UCEPHAN			
B BRAUN	100MG/ML;100MG/ML	N019530 001	Dec 23, 1987

SODIUM BICARBONATE

INJECTABLE; INJECTION

SODIUM BICARBONATE			
HOSPIRA	0.9MEQ/ML	A077394 001	Nov 09, 2005
	1MEQ/ML	A077394 002	Nov 09, 2005
SODIUM BICARBONATE IN PLASTIC CONTAINER			
+ ABBOTT	0.9MEQ/ML **	N019443 001	Jun 03, 1986
+	1MEQ/ML **	N019443 002	Jun 03, 1986

SODIUM BICARBONATE; TARTARIC ACID

GRANULE, EFFERVESCENT; ORAL

BAROS			
MALLINCKRODT INC	460MG/GM;420MG/GM	N018509 001	Aug 07, 1985

SODIUM CHLORIDE

AEROSOL, METERED; INHALATION

BRONCHO SALINE			
+ BLAIREX	0.9%	N019912 001	Sep 03, 1992

INJECTABLE; INJECTION

SODIUM CHLORIDE			
ABBOTT	20GM/100ML	N017013 001	
B BRAUN	20GM/100ML	N017038 001	
SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER			
B BRAUN	450MG/100ML	N018184 001	
MILES	450MG/100ML	N018503 001	
SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER			
+ LIEBEL-FLARSHEIM	450MG/50ML (9MG/ML)	N021569 001	Jul 27, 2006
SODIUM CHLORIDE 23.4% IN PLASTIC CONTAINER			
+ ABRAXIS PHARM	234MG/ML **	N019329 001	Apr 22, 1987
SOLUTION; INJECTION			
BACTERIOSTATIC SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER			
ABRAXIS PHARM	90MG/10ML (9MG/ML)	A088909 002	May 15, 1985
	270MG/30ML (9MG/ML)	A088909 001	Feb 07, 1985

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

SODIUM CHLORIDE

SOLUTION; INJECTION

SODIUM CHLORIDE 0.9%				
	HIKMA	90MG/10ML (9MG/ML)	A201850 001	Jan 20, 2012
SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER				
	ABBOTT	90MG/10ML (9MG/ML)	N019218 001	Jul 13, 1984
		180MG/20ML (9MG/ML)	N019218 002	Apr 30, 1985
+	HOSPIRA	18MG/2ML (9MG/ML)	N018803 004	Jan 22, 2015
+		27MG/3ML (9MG/ML)	N018803 005	Jan 22, 2015
+		45MG/5ML (9MG/ML)	N018803 006	Jan 22, 2015
+	ICU MEDICAL INC	45MG/5ML (9MG/ML)	N019217 002	Nov 18, 1998
+		90MG/10ML (9MG/ML)	N019217 003	Nov 18, 1998
+		360MG/40ML (9MG/ML)	N019217 004	Apr 01, 1999
	JUBILANT CADISTA	27MG/3ML (9MG/ML)	A203352 002	May 18, 2016
		90MG/10ML (9MG/ML)	A203352 001	May 18, 2016
	MILES	450MG/50ML (9MG/ML)	N018502 002	
		900MG/100ML	N018502 001	
SODIUM CHLORIDE 0.9% IN PLASTIC THERMOJECT KIT FOR CARDIAC OUTPUT USE				
+	ICU MEDICAL INC	90MG/10ML (9MG/ML)	N019217 001	Jul 13, 1984

SOLUTION; INTRAVENOUS

SODIUM CHLORIDE 14.6%				
+	HOSPIRA	50MEQ/20ML (2.5MEQ/ML) **	N018897 001	Jul 20, 1984

SOLUTION; IRRIGATION

SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER				
	BAXTER HLTHCARE	450MG/100ML	N017864 001	
		450MG/100ML	N018497 001	Feb 19, 1982
	HOSPIRA	450MG/100ML	N017670 001	
		450MG/100ML	N018380 001	
SODIUM CHLORIDE IN PLASTIC CONTAINER				
	MILES	900MG/100ML	N018247 001	

SODIUM CHROMATE CR-51

INJECTABLE; INJECTION

CHROMITOPE SODIUM				
	BRACCO	2mCi/VIAL	N013993 002	
		200uCi/ML	N013993 001	
SODIUM CHROMATE CR 51				
	CURIUM	100uCi/ML	N016708 001	

SODIUM FLUORIDE F-18

INJECTABLE; INTRAVENOUS

FLUORINE F-18				
+	GE HEALTHCARE	2mCi/ML **	N017042 001	
SODIUM FLUORIDE F 18				
+	NIH NCI DCTD	10-200mCi/ML **	N022494 001	Jan 26, 2011
SODIUM FLUORIDE F-18				
	DECATUR	10-200mCi/ML	A204464 001	Oct 21, 2014
	SHERTECH LABS LLC	10-200mCi/ML	A204315 001	Sep 22, 2014
	SOFIE	10-200mCi/ML	A203544 001	Dec 26, 2012
	UCSF RODIOPHARM	10-200mCi/ML	A204437 001	Mar 13, 2014
	UIHC PET IMAGING	10-200mCi/ML	A204462 001	Nov 17, 2015
	UNIV TX MD ANDERSON	10-200mCi/ML	A203247 001	Dec 23, 2013

SODIUM FLUORIDE; TRICLOSAN

PASTE; DENTAL

COLGATE TOTAL				
+	COLGATE PALMOLIVE	0.24%; 0.3%	N020231 001	Jul 11, 1997

SODIUM IODIDE I-123

CAPSULE; ORAL

SODIUM IODIDE I 123				
	CARDINAL HEALTH 418	400uCi	N018671 003	May 27, 1982
	GE HEALTHCARE	100uCi	N017630 001	
		200uCi	N017630 003	Jan 08, 1993

SOLUTION; ORAL

SODIUM IODIDE I 123				
	GE HEALTHCARE	2mCi/ML **	N017630 002	

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

SODIUM IODIDE I-131

CAPSULE; ORAL

IODOTOPE

BRACCO	1-130mCi	N010929 001
	1-150mCi	N010929 003

SODIUM IODIDE I 131

CIS	50uCi	N017316 001
	100uCi	N017316 002
CURIUM	0.8-100mCi	N016515 002
+	0.8-100mCi	N016517 001
	15-100uCi	N016517 002
JUBILANT	2-200mCi	N021305 004 Nov 18, 2004

SOLUTION; ORAL

HICON

JUBILANT	1-250mCi/0.25ML	N021305 002 Jan 24, 2003
	1-500mCi/0.5ML	N021305 003 Jan 24, 2003
	1-1000mCi/ML	N021305 005 Apr 04, 2006

IODOTOPE

BRACCO	7-106mCi/BOT	N010929 002
--------	--------------	-------------

SODIUM IODIDE I 131

CIS	50mCi/ML	N017315 001
+	CURIUM	3.5-150mCi/VIAL N016515 001

SODIUM LACTATE

INJECTABLE; INJECTION

SODIUM LACTATE 0.167 MOLAR IN PLASTIC CONTAINER

B BRAUN	1.87GM/100ML	N018186 001
BAXTER HLTHCARE	1.87GM/100ML	N016692 001
HOSPIRA	1.87GM/100ML	N018249 001

SODIUM LACTATE 1/6 MOLAR IN PLASTIC CONTAINER

B BRAUN	1.87GM/100ML	N020004 001 Apr 21, 1992
---------	--------------	--------------------------

SODIUM LACTATE IN PLASTIC CONTAINER

+	HOSPIRA	5MEQ/ML N018947 001 Sep 05, 1984
---	---------	----------------------------------

SODIUM MONOFLUOROPHOSPHATE

GEL; DENTAL

EXTRA-STRENGTH AIM

CHESEBROUGH PONDS	1.2%	N019518 002 Aug 06, 1986
-------------------	------	--------------------------

PASTE; DENTAL

EXTRA-STRENGTH AIM

CHESEBROUGH PONDS	1.2%	N019518 001 Jun 03, 1987
-------------------	------	--------------------------

SODIUM NITROPRUSSIDE

INJECTABLE; INJECTION

NIPRIDE

ROCHE	50MG/VIAL	N017546 001
-------	-----------	-------------

NITROPRESS

ABBOTT	50MG/VIAL	A071555 001 Nov 16, 1987
+	ABBVIE	50MG/VIAL ** N018450 001
HOSPIRA	50MG/VIAL	A070566 001 Jun 09, 1986
VPNA	25MG/ML	A071961 001 Aug 01, 1988

SODIUM NITROPRUSSIDE

ABRAXIS PHARM	50MG/VIAL	A070031 001 Jan 17, 1985
AMPHASTAR PHARMS INC	25MG/ML	A209832 001 Dec 18, 2017
AVET LIFESCIENCES	25MG/ML	A208923 001 Nov 08, 2022
+	BAXTER HLTHCARE	50MG/VIAL ** N018581 001 Jul 28, 1982
CHARTWELL RX	25MG/ML	A209834 001 Jun 26, 2018
CIPLA	25MG/ML	A210855 001 Jul 16, 2018
EPIC PHARMA LLC	25MG/ML	A208635 001 May 04, 2017
EUGIA PHARMA	25MG/ML	A211934 001 Dec 10, 2020
SUN PHARM	25MG/ML	A210467 001 Nov 26, 2018
TEVA PARENTERAL	25MG/ML	A073465 001 Mar 30, 1992

SOLUTION; INTRAVENOUS

NIPRIDE RTU IN SODIUM CHLORIDE 0.9%

+	EXELA PHARMA	10MG/50ML (0.2MG/ML) ** N209387 002 Dec 07, 2017
---	--------------	--

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

SODIUM OXYBATE

SOLUTION;ORAL

SODIUM OXYBATE

HIKMA

0.5GM/ML

A202090 001 Jan 17, 2017

SODIUM PHENYLBUTYRATE

TABLET;ORAL

SODIUM PHENYLBUTYRATE

ALVOGEN

500MG

A090910 001 Nov 18, 2011

SODIUM PHENYLBUTYRATE; TAURURSODIOL

FOR SUSPENSION;ORAL

RELYVRIO

+ AMYLYX

3GM/PACKET; 1GM/PACKET

N216660 001 Sep 29, 2022

SODIUM PHOSPHATE P-32

SOLUTION;INJECTION, ORAL

PHOSPHOTOPE

BRACCO

1-8mCi/VIAL

N010927 001

SODIUM PHOSPHATE P 32

MALLINCKRODT

0.67mCi/ML

N011777 001

1.5mCi/VIAL

N011777 002

SODIUM PHOSPHATE, DIBASIC, ANHYDROUS; SODIUM PHOSPHATE, MONOBASIC, MONOHYDRATE

TABLET;ORAL

OSMOPREP

+ SALIX PHARMS

0.398GM;1.102GM **

N021892 001 Mar 16, 2006

VISICOL

SALIX PHARMS

0.398GM;1.102GM

N021097 001 Sep 21, 2000

SODIUM POLYSTYRENE SULFONATE

POWDER;ORAL, RECTAL

KAYEXALATE

+ CONCORDIA

453.6GM/BOT **

N011287 001

SODIUM POLYSTYRENE SULFONATE

CITRUSPHRMA

454GM/BOT

A040909 001 Dec 03, 2008

+ PAI HOLDINGS PHARM

453.6GM/BOT

A088786 001 Sep 11, 1984

SUSPENSION;ORAL, RECTAL

SODIUM POLYSTYRENE SULFONATE

ANI PHARMS

15GM/60ML

A090590 001 May 13, 2011

HIKMA

15GM/60ML

A089049 001 Nov 17, 1986

MORTON GROVE

15GM/60ML

A088717 001 Sep 11, 1984

ROXANE

15GM/60ML

A088453 001 Nov 17, 1983

SODIUM SUCCINATE

INJECTABLE;INJECTION

SODIUM SUCCINATE

ELKINS SINN

30%

A080516 001

SODIUM TETRADECYL SULFATE

INJECTABLE;INJECTION

SOTRADECOL

+ ELKINS SINN

1% **

N005970 004

+

3% **

N005970 005

SODIUM THIOSULFATE

INJECTABLE;INJECTION

SODIUM THIOSULFATE

+ US ARMY

250MG/ML

N020166 001 Feb 14, 1992

SOFOSBUVIR

TABLET;ORAL

SOFOSBUVIR

TEVA PHARMS USA INC

400MG

A211353 001 Jan 27, 2022

SOLIFENACIN SUCCINATE

TABLET;ORAL

SOLIFENACIN SUCCINATE

ACCORD HLTHCARE

5MG

A207477 001 Jan 04, 2022

10MG

A207477 002 Jan 04, 2022

AJANTA PHARMA LTD

5MG

A205483 001 May 20, 2019

10MG

A205483 002 May 20, 2019

AMNEAL PHARMS CO

5MG

A209719 001 May 20, 2019

10MG

A209719 002 May 20, 2019

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

SOLIFENACIN SUCCINATE

TABLET; ORAL

SOLIFENACIN SUCCINATE

BRECKENRIDGE	5MG	A209818 001	May 20, 2019
	10MG	A209818 002	May 20, 2019
CIPLA	5MG	A209839 001	May 20, 2019
	10MG	A209839 002	May 20, 2019
LANNETT CO INC	5MG	A211622 001	Jun 06, 2023
	10MG	A211622 002	Jun 06, 2023
SUNSHINE	5MG	A213346 001	Apr 13, 2020
	10MG	A213346 002	Apr 13, 2020
ZYDUS PHARMS	5MG	A207721 001	Oct 19, 2020
	10MG	A207721 002	Oct 19, 2020

SORBITOL

SOLUTION; IRRIGATION

SORBITOL 3% IN PLASTIC CONTAINER

BAXTER HLTHCARE	3GM/100ML	N018512 001	May 27, 1982
-----------------	-----------	-------------	--------------

SOTALOL HYDROCHLORIDE

TABLET; ORAL

BETAPACE

LEGACY PHARMA	320MG	N019865 004	Oct 30, 1992
---------------	-------	-------------	--------------

BETAPACE AF

LEGACY PHARMA	40MG	N021151 006	Apr 02, 2003
---------------	------	-------------	--------------

	60MG	N021151 007	Apr 02, 2003
--	------	-------------	--------------

	100MG	N021151 005	Mar 14, 2003
--	-------	-------------	--------------

SOTALOL HYDROCHLORIDE

ESJAY PHARMA	80MG	A075237 001	May 01, 2000
--------------	------	-------------	--------------

	120MG	A075237 002	May 01, 2000
--	-------	-------------	--------------

	160MG	A075237 003	May 01, 2000
--	-------	-------------	--------------

	240MG	A075237 004	May 01, 2000
--	-------	-------------	--------------

IMPAX PHARMS	80MG	A075663 001	Nov 07, 2000
--------------	------	-------------	--------------

	120MG	A075663 002	Nov 07, 2000
--	-------	-------------	--------------

	160MG	A075663 003	Nov 07, 2000
--	-------	-------------	--------------

	240MG	A075663 004	Nov 07, 2000
--	-------	-------------	--------------

RISING	80MG	A075725 001	Dec 19, 2000
--------	------	-------------	--------------

	120MG	A075725 002	Dec 19, 2000
--	-------	-------------	--------------

	160MG	A075725 003	Dec 19, 2000
--	-------	-------------	--------------

	240MG	A075725 004	Dec 19, 2000
--	-------	-------------	--------------

SUN PHARM INDUSTRIES	80MG	A075515 001	Oct 15, 2001
----------------------	------	-------------	--------------

	80MG	A076576 001	Apr 08, 2004
--	------	-------------	--------------

	120MG	A075515 004	Oct 15, 2001
--	-------	-------------	--------------

	120MG	A076576 002	Apr 08, 2004
--	-------	-------------	--------------

	160MG	A075515 002	Oct 15, 2001
--	-------	-------------	--------------

	160MG	A076576 003	Apr 08, 2004
--	-------	-------------	--------------

	240MG	A075515 003	Oct 15, 2001
--	-------	-------------	--------------

TEVA	80MG	A076883 001	Jul 26, 2004
------	------	-------------	--------------

	120MG	A076883 002	Jul 26, 2004
--	-------	-------------	--------------

	160MG	A076883 003	Jul 26, 2004
--	-------	-------------	--------------

UPSHER SMITH LABS	80MG	A075366 001	May 01, 2000
-------------------	------	-------------	--------------

	120MG	A075366 002	May 01, 2000
--	-------	-------------	--------------

	160MG	A075366 003	May 01, 2000
--	-------	-------------	--------------

	240MG	A075366 004	May 01, 2000
--	-------	-------------	--------------

WATSON LABS	80MG	A075238 001	Jul 13, 2000
-------------	------	-------------	--------------

	120MG	A075238 002	Jul 13, 2000
--	-------	-------------	--------------

	160MG	A075238 003	Jul 13, 2000
--	-------	-------------	--------------

	240MG	A075238 004	Jul 13, 2000
--	-------	-------------	--------------

SOYBEAN OIL

INJECTABLE; INJECTION

INTRALIPID 10%

+ FRESENIUS

	10%	N017643 001	
--	-----	-------------	--

LIPOSYN III 10%

HOSPIRA	10%	N018969 001	Sep 24, 1984
---------	-----	-------------	--------------

LIPOSYN III 20%

HOSPIRA	20%	N018970 001	Sep 25, 1984
---------	-----	-------------	--------------

LIPOSYN III 30%

HOSPIRA	30%	N020181 001	Jan 13, 1998
---------	-----	-------------	--------------

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

SOYBEAN OIL

INJECTABLE; INJECTION

NUTRILIPID 10%

+ B BRAUN

10%

N019531 001 May 28, 1993

SOYACAL 10%

ALPHA THERA

10%

N018465 001 Jun 29, 1983

SOYACAL 20%

ALPHA THERA

20%

N018786 001 Jun 29, 1983

TRAVAMULSION 10%

BAXTER HLTHCARE

10%

N018660 001 Feb 26, 1982

TRAVAMULSION 20%

BAXTER HLTHCARE

20%

N018758 001 Feb 15, 1983

SPARFLOXACIN

TABLET; ORAL

ZAGAM

MYLAN

200MG

N020677 001 Dec 19, 1996

SPECTINOMYCIN HYDROCHLORIDE

INJECTABLE; INJECTION

TROBICIN

PFIZER

EQ 2GM BASE/VIAL

N050347 001

EQ 4GM BASE/VIAL

N050347 002

SPIRAPRIL HYDROCHLORIDE

TABLET; ORAL

RENORMAX

SCHERING

3MG

N020240 001 Dec 29, 1994

6MG

N020240 002 Dec 29, 1994

12MG

N020240 003 Dec 29, 1994

24MG

N020240 004 Dec 29, 1994

SPIRONOLACTONE

TABLET; ORAL

SPIRONOLACTONE

ACTAVIS ELIZABETH

25MG

A040353 003 Mar 15, 2006

50MG

A040353 001 Jul 29, 1999

100MG

A040353 002 Jul 29, 1999

ASCOT

25MG

A087687 001 Oct 20, 1982

CHARTWELL RX

25MG

A086809 001

IVAX PHARMS

25MG

A087108 001

LEDERLE

25MG

A087634 001

MUTUAL PHARM

25MG

A087265 001

NORVIUM BIOSCIENCE

25MG

A087086 001

PUREPAC PHARM

25MG

A087998 001 Oct 14, 1983

25MG

A088053 001 Aug 25, 1983

SUPERPHARM

25MG

A089364 001 Nov 07, 1986

UPSHER SMITH

25MG

A087554 001

VANGARD

25MG

A087648 001 Feb 01, 1982

WARNER CHILCOTT

25MG

A087952 001 Nov 18, 1982

WATSON LABS

25MG

A086898 002 Mar 02, 1982

25MG

A087078 001

STANOZOLOL

TABLET; ORAL

WINSTROL

+ LUNDBECK INC

2MG

N012885 001 May 14, 1984

STAVUDINE

CAPSULE; ORAL

STAVUDINE

AUROBINDO PHARMA

15MG

A077672 003 Dec 29, 2008

20MG

A077672 004 Dec 29, 2008

30MG

A077672 001 Dec 29, 2008

40MG

A077672 002 Dec 29, 2008

HETERO LABS LTD III

15MG

A078957 001 Dec 29, 2008

20MG

A078957 002 Dec 29, 2008

30MG

A078957 003 Dec 29, 2008

40MG

A078957 004 Dec 29, 2008

NORVIUM BIOSCIENCE

15MG

A079069 001 Dec 29, 2008

20MG

A079069 002 Dec 29, 2008

30MG

A078775 001 Jan 05, 2009

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

STAVUDINE

CAPSULE; ORAL

STAVUDINE

30MG	A079069	003	Dec 29, 2008
40MG	A078775	002	Jan 05, 2009
40MG	A079069	004	Dec 29, 2008

ZERIT

BRISTOL

5MG N020412 001 Jun 24, 1994

+ 15MG ** N020412 002 Jun 24, 1994

+ 20MG ** N020412 003 Jun 24, 1994

+ 30MG ** N020412 004 Jun 24, 1994

+ 40MG ** N020412 005 Jun 24, 1994

CAPSULE, EXTENDED RELEASE; ORAL

ZERIT XR

BRISTOL MYERS SQUIBB

37.5MG N021453 001 Dec 31, 2002

50MG N021453 002 Dec 31, 2002

75MG N021453 003 Dec 31, 2002

100MG N021453 004 Dec 31, 2002

FOR SOLUTION; ORAL

STAVUDINE

AUROBINDO PHARMA

1MG/ML A077774 001 Dec 29, 2008

CIPLA LTD

1MG/ML A078030 001 Mar 20, 2009

ZERIT

+ BRISTOL

1MG/ML ** N020413 001 Sep 06, 1996

STERILE WATER FOR INJECTION

LIQUID; N/A

BACTERIOSTATIC WATER FOR INJECTION IN PLASTIC CONTAINER

ABRAXIS PHARM

100% A089099 001 Dec 29, 1987

100% A089100 001 Dec 29, 1987

STERILE WATER FOR INJECTION

HIKMA

100% (20ML) A206369 002 Sep 02, 2015

+ HOSPIRA 100% (1ML) ** N018801 001 Oct 27, 1982

+ 100% (5.2ML) ** N018801 006 May 02, 2023

STERILE WATER FOR INJECTION IN PLASTIC CONTAINER

B BRAUN

100% N019077 001 Mar 02, 1984

STERILE WATER FOR IRRIGATION

LIQUID; IRRIGATION

STERILE WATER IN PLASTIC CONTAINER

MILES

100% N018246 001

STREPTOMYCIN SULFATE

INJECTABLE; INJECTION

STREPTOMYCIN SULFATE

COPANOS

EQ 500MG BASE/ML A060684 001

LILLY

EQ 1GM BASE/VIAL A060107 001

EQ 1GM BASE/2ML A060404 001

EQ 5GM BASE/VIAL A060107 002

PFIZER

EQ 1GM BASE/VIAL ** A060076 001

EQ 1GM BASE/2.5ML A060111 001

EQ 5GM BASE/VIAL ** A060076 002

SUCCINYLCHOLINE CHLORIDE

INJECTABLE; INJECTION

ANECTINE

SANDOZ

50MG/ML N008453 003

500MG/VIAL N008453 001

1GM/VIAL N008453 004

QUELICIN PRESERVATIVE FREE

+ HOSPIRA

20MG/ML ** N008845 001

+ 50MG/ML ** N008845 002

+ 100MG/ML ** N008845 004

SUCCINYLCHOLINE CHLORIDE

ACCORD HLTHCARE

20MG/ML A213705 001 May 20, 2020

AMPHASTAR PHARMS INC

20MG/ML A213432 001 Jun 08, 2020

BRECKENRIDGE

20MG/ML A212638 001 Oct 09, 2019

EUGIA PHARMA

20MG/ML A217808 001 Oct 16, 2023

INTL MEDICATION

100MG/VIAL A085400 001 Feb 04, 1982

MANKIND PHARMA

20MG/ML A216127 001 Feb 02, 2023

NEXUS

20MG/ML A213552 001 Oct 27, 2020

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

SUCCINYLCHOLINE CHLORIDE

INJECTABLE; INJECTION

SUCCINYLCHOLINE CHLORIDE

ORGANON USA INC

20MG/ML

A080997 001

SUCOSTRIN

+ APOTHECON

20MG/ML

N008847 001

+

100MG/ML

N008847 003

SUCRALFATE

SUSPENSION; ORAL

SUCRALFATE

HIKMA

1GM/10ML

A212769 001 Jul 23, 2024

SUFENTANIL CITRATE

INJECTABLE; INJECTION

SUFENTANIL CITRATE

WATSON LABS

EQ 0.05MG BASE/ML

A074406 001 Dec 15, 1995

TABLET; SUBLINGUAL

DSUVIA

+ VERTICAL PHARMS

EQ 0.03MG BASE

N209128 001 Nov 02, 2018

SUGAMMADEX SODIUM

SOLUTION; INTRAVENOUS

SUGAMMADEX SODIUM

ZYDUS PHARMS

EQ 200MG BASE/2ML (EQ 100MG BASE/ML)

A214290 001 Oct 04, 2023

EQ 500MG BASE/5ML (EQ 100MG BASE/ML)

A214290 002 Oct 04, 2023

SULFACETAMIDE SODIUM

OINTMENT; OPHTHALMIC

BLEPH-10

ALLERGAN

10%

A084015 001

CETAMIDE

ALCON

10%

A080021 001

SODIUM SULAMYD

+ SCHERING

10% **

N005963 002

SULFAIR 10

PHARMAFAIR

10%

A088000 001 Dec 22, 1982

SOLUTION/DROPS; OPHTHALMIC

BLEPH-10

+ ALLERGAN

10%

A080028 001

BLEPH-30

ALLERGAN

30%

A080028 002

ISOPTO CETAMIDE

ALCON

15%

A080020 002

OCUSULF-10

MIZA PHARMS USA

10%

A080660 001

OCUSULF-30

MIZA PHARMS USA

30%

A080660 002

SODIUM SULAMYD

+ SCHERING

10% **

N005963 001

+

30% **

N005963 003

SODIUM SULFACETAMIDE

EPIC PHARMA LLC

10%

A083021 001

15%

A083021 002

30%

A083021 003

SOLA BARNES HIND

10%

A084143 001

10%

A084145 001

30%

A084146 001

30%

A084147 001

SULF-10

NOVARTIS

10%

A080025 001

SULF-15

NOVARTIS

15%

A089047 001 Oct 31, 1995

SULFACEL-15

OPTOPICS

15%

A080024 001

SULFACETAMIDE SODIUM

ALCON PHARMS LTD

30%

A089068 001 May 05, 1987

EPIC PHARMA LLC

30%

A040216 001 May 25, 1999

PHARMAFAIR

10%

A088947 001 May 17, 1985

RENOVA PHARMS

10%

A040215 001 May 25, 1999

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

SULFACETAMIDE SODIUM

SOLUTION/DROPS;OPHTHALMIC

SULFAIR 10

PHARMAFAIR 10% A087949 001 Dec 13, 1982

SULFAIR FORTE

PHARMAFAIR 30% A088385 001 Oct 13, 1983

SULFAIR-15

PHARMAFAIR 15% A088186 001 May 25, 1983

SULTEN-10

BAUSCH AND LOMB 10% A087818 001 Feb 03, 1983

SULFACYTINE

TABLET;ORAL

RENOQUID

GLENWOOD 250MG N017569 001

SULFADIAZINE

TABLET;ORAL

SULFADIAZINE

ABBVIE 300MG N004125 005

CHARTWELL MOLECULAR 500MG A080084 001

EVERYLIFE 500MG A080088 001

+ IMPAX LABS 500MG A080081 001

LEDERLE 500MG N004054 001

+ LILLY 500MG N004122 002

SULFADIAZINE SODIUM

INJECTABLE; INJECTION

SULFADIAZINE SODIUM

LEDERLE 250MG/ML N004054 002

SULFADIAZINE; SULFAMERAZINE

SUSPENSION;ORAL

SULFONAMIDES DUPLEX

LILLY 250MG/5ML;250MG/5ML N006317 007

SULFAMETER

TABLET;ORAL

SULLA

BAYER HLTHCARE 500MG N016000 002

SULFAMETHIZOLE

TABLET;ORAL

MICROSUL

FOREST PHARMS 1GM A086012 001

PROKLAR

FOREST PHARMS 500MG A080273 001

THIOSULFIL

WYETH AYERST 250MG N008565 001

500MG N008565 004

SULFAMETHOXAZOLE

SUSPENSION;ORAL

GANTANOL

ROCHE 500MG/5ML N013664 002

TABLET;ORAL

GANTANOL

ROCHE 500MG N012715 002

GANTANOL-DS

ROCHE 1GM N012715 003

SULFAMETHOXAZOLE

ASCOT 500MG A087662 001 Oct 20, 1982

BARR 500MG A087189 001 Jul 25, 1983

HEATHER 500MG A086163 001

RISING 500MG A085844 001

WATSON LABS 500MG A085053 001

1GM A086000 001

UROBAK

SHIONOGI 500MG A087307 001

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

SULFAMETHOXAZOLE; TRIMETHOPRIM

INJECTABLE; INJECTION

BACTRIM

+ SUN PHARM INDS INC 80MG/ML;16MG/ML ** N018374 001

SEPTRA

MONARCH PHARMS 80MG/ML;16MG/ML ** N018452 001

SULFAMETHOXAZOLE AND TRIMETHOPRIM

ABRAXIS PHARM 80MG/ML;16MG/ML A070223 001 Dec 29, 1987

BEDFORD 80MG/ML;16MG/ML A072383 001 Apr 29, 1992

HIKMA 80MG/ML;16MG/ML A070627 001 Dec 29, 1987

80MG/ML;16MG/ML A070628 001 Dec 29, 1987

HOSPIRA 80MG/ML;16MG/ML A073199 001 Sep 11, 1992

WATSON LABS 80MG/ML;16MG/ML A071556 001 Dec 29, 1987

SUSPENSION; ORAL

BACTRIM

+ SUN PHARM INDUSTRIES 200MG/5ML;40MG/5ML ** N017560 001

BACTRIM PEDIATRIC

SUN PHARM INDUSTRIES 200MG/5ML;40MG/5ML ** N017560 002

SEPTRA

MONARCH PHARMS 200MG/5ML;40MG/5ML ** N017598 001

SEPTRA GRAPE

MONARCH PHARMS 200MG/5ML;40MG/5ML ** N017598 002 Feb 12, 1986

SULFAMETHOXAZOLE AND TRIMETHOPRIM

ANI PHARMS 200MG/5ML;40MG/5ML A070028 001 Jun 02, 1987

HIKMA 200MG/5ML;40MG/5ML A074650 001 Dec 29, 1997

LUPIN LTD 200MG/5ML;40MG/5ML A212699 001 Jan 05, 2021

TEVA 200MG/5ML;40MG/5ML N018812 001 Jan 28, 1983

200MG/5ML;40MG/5ML N018812 002 Jun 10, 1983

SULFATRIM

PHARM ASSOC 200MG/5ML;40MG/5ML N018615 002 Jan 07, 1983

SULMEPRIM

USL PHARMA 200MG/5ML;40MG/5ML A070063 001 Aug 01, 1986

SULMEPRIM PEDIATRIC

USL PHARMA 200MG/5ML;40MG/5ML A070064 001 Aug 01, 1986

TRIMETH/SULFA

ALPHARMA US PHARMS 200MG/5ML;40MG/5ML A072289 001 May 23, 1988

200MG/5ML;40MG/5ML A072398 001 May 23, 1988

NASKA 200MG/5ML;40MG/5ML A072399 001 May 23, 1988

TABLET; ORAL

COTRIM

TEVA 400MG;80MG A070034 001 May 16, 1985

COTRIM D.S.

TEVA 800MG;160MG A070048 001 Mar 18, 1985

SULFAMETHOPRIM

NOVEL LABS INC 400MG;80MG A070022 001 Feb 15, 1985

SULFAMETHOPRIM-DS

NOVEL LABS INC 800MG;160MG A070032 001 Feb 15, 1985

SULFAMETHOXAZOLE AND TRIMETHOPRIM

AIPING PHARM INC 400MG;80MG A070889 001 Nov 13, 1986

800MG;160MG A070890 001 Nov 13, 1986

FOSUN PHARMA 400MG;80MG N018598 003 May 19, 1982

HEATHER 400MG;80MG N018946 001 Aug 10, 1984

800MG;160MG N018946 002 Aug 10, 1984

INTERPHARM 400MG;80MG A071299 001 Oct 27, 1987

800MG;160MG A071300 001 Oct 27, 1987

MARTEC USA LLC 400MG;80MG A072408 001 Dec 07, 1988

MUTUAL PHARM 400MG;80MG A070006 001 Nov 14, 1984

PLIVA 400MG;80MG A070215 001 Sep 10, 1985

800MG;160MG A070216 001 Sep 10, 1985

ROXANE 400MG;80MG A072768 001 Aug 30, 1991

TEVA 400MG;80MG N018242 001

800MG;160MG N018242 002

USL PHARMA 400MG;80MG A070203 001 Nov 08, 1985

800MG;160MG A070204 001 Nov 08, 1985

WATSON LABS 400MG;80MG A070002 001 Nov 07, 1984

400MG;80MG N018852 001 May 09, 1983

800MG;160MG A070000 001 Nov 07, 1984

SULFAMETHOXAZOLE AND TRIMETHOPRIM DOUBLE STRENGTH

FOSUN PHARMA 800MG;160MG N018598 004 May 19, 1982

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

SULFAMETHOXAZOLE; TRIMETHOPRIM

TABLET; ORAL

SULFAMETHOXAZOLE AND TRIMETHOPRIM DOUBLE STRENGTH

HERITAGE PHARMA AVET	800MG;160MG	A070037	001	Jun 02, 1987
MARTEC USA LLC	800MG;160MG	A072417	001	Dec 07, 1988
MUTUAL PHARM	800MG;160MG	A070007	001	Nov 14, 1984
ROXANE	800MG;160MG	A072769	001	Aug 30, 1991
WATSON LABS	800MG;160MG	N018854	001	May 09, 1983

SULFAMETHOXAZOLE AND TRIMETHOPRIM SINGLE STRENGTH

HERITAGE PHARMA AVET	400MG;80MG	A070030	001	Jun 02, 1987
SULFATRIM-DS				
SUPERPHARM	800MG;160MG	A070066	001	Jun 24, 1985
SULFATRIM-SS				
SUPERPHARM	400MG;80MG	A070065	002	Jun 24, 1985
UROPLUS DS				
SHIONOGI	800MG;160MG	A071816	001	Sep 28, 1987
UROPLUS SS				
SHIONOGI	400MG;80MG	A071815	001	Sep 28, 1987

SULFANILAMIDE

CREAM; VAGINAL

AVC				
+ NORVIUM BIOSCIENCE	15% **	N006530	003	Jan 27, 1987
SULFANILAMIDE				
COSETTE	15%	A088718	001	Sep 19, 1985

SUPPOSITORY; VAGINAL

AVC				
NORVIUM BIOSCIENCE	1.05GM	N006530	004	Jan 27, 1987

SULFAPHENAZOLE

SUSPENSION; ORAL

SULFABID				
PHARM RES ASSOC	500MG/5ML	N013093	001	

TABLET; ORAL

SULFABID				
PURDUE FREDERICK	500MG	N013092	002	

SULFAPYRIDINE

TABLET; ORAL

SULFAPYRIDINE				
LILLY	500MG	N000159	001	

SULFASALAZINE

SUSPENSION; ORAL

AZULFIDINE				
PHARMACIA AND UPJOHN	250MG/5ML	N018605	001	

TABLET; ORAL

S.A.S.-500				
SOLVAY	500MG	A083450	001	
SULFASALAZINE				
EPIC PHARMA LLC	500MG	A086184	001	
SUN PHARM INDUSTRIES	500MG	A089590	001	Oct 19, 1987
SUPERPHARM	500MG	A089339	001	Oct 26, 1987
WATSON LABS	500MG	A084964	001	
	500MG	A087197	001	

TABLET, DELAYED RELEASE; ORAL

SULFASALAZINE				
WATSON LABS	500MG	A088052	001	May 24, 1983

SULFINPYRAZONE

CAPSULE; ORAL

ANTURANE				
+ NOVARTIS	200MG **	N011556	004	

SULFINPYRAZONE

BARR	200MG	A087666	001	Sep 17, 1982
IVAX PHARMS	200MG	A087770	001	Nov 19, 1982
PAR PHARM	200MG	A088934	001	Sep 06, 1985
VANGARD	200MG	A088666	001	Feb 17, 1984

TABLET; ORAL

ANTURANE				
NOVARTIS	100MG **	N011556	003	

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

SULFINPYRAZONE

TABLET; ORAL

SULFINPYRAZONE

BARR	100MG	A087665 001	Sep 17, 1982
IVAX PHARMS	100MG	A087769 001	Jun 01, 1982
PAR PHARM	100MG	A088933 001	Sep 06, 1985
WATSON LABS	100MG	A087667 001	May 26, 1982

SULFISOXAZOLE

TABLET; ORAL

GANTRISIN

ROCHE	500MG	N006525 001	
-------	-------	-------------	--

SOSOL

MK LABS	500MG	A080036 001	
---------	-------	-------------	--

SOXAZOLE

ALRA	500MG	A080366 001	
------	-------	-------------	--

SULFALAR

PARKE DAVIS	500MG	A084955 001	
-------------	-------	-------------	--

SULFISOXAZOLE

ANI PHARMS	500MG	A080142 001	
------------	-------	-------------	--

BARR	500MG	A084031 001	
------	-------	-------------	--

HEATHER	500MG	A080189 001	
---------	-------	-------------	--

IMPAX LABS	500MG	A080109 001	
------------	-------	-------------	--

LANNETT	500MG	A080085 001	
---------	-------	-------------	--

LEDERLE	500MG	A087649 001	
---------	-------	-------------	--

PHARMERAL	500MG	A084385 001	
-----------	-------	-------------	--

PUREPAC PHARM	500MG	A080087 001	
---------------	-------	-------------	--

RISING	500MG	A085628 001	
--------	-------	-------------	--

ROXANE	500MG	A080082 001	
--------	-------	-------------	--

VALEANT PHARM INTL	500MG	A080268 002	
--------------------	-------	-------------	--

VITARINE	500MG	A087332 001	
----------	-------	-------------	--

WATSON LABS	500MG	A085534 001	
-------------	-------	-------------	--

WEST WARD	500MG	A080379 001	
-----------	-------	-------------	--

SULSOXIN

SOLVAY	500MG	A080040 001	
--------	-------	-------------	--

SULFISOXAZOLE ACETYL

EMULSION; ORAL

LIPO GANTRISIN

ROCHE	EQ 1GM BASE/5ML	N009182 009	
-------	-----------------	-------------	--

SUSPENSION; ORAL

GANTRISIN PEDIATRIC

ROCHE	EQ 500MG BASE/5ML	N009182 004	
-------	-------------------	-------------	--

SYRUP; ORAL

GANTRISIN

ROCHE	EQ 500MG BASE/5ML	N009182 002	
-------	-------------------	-------------	--

SULFISOXAZOLE DIOLAMINE

INJECTABLE; INJECTION

GANTRISIN

ROCHE	EQ 400MG BASE/ML	N006917 001	
-------	------------------	-------------	--

OINTMENT; OPHTHALMIC

GANTRISIN

ROCHE	EQ 4% BASE	N008414 002	
-------	------------	-------------	--

SOLUTION/DROPS; OPHTHALMIC

GANTRISIN

ROCHE	EQ 4% BASE	N007757 002	
-------	------------	-------------	--

SULFISOXAZOLE DIOLAMINE

SOLA BARNES HIND	EQ 4% BASE	A084148 001	
------------------	------------	-------------	--

SULFOXONE SODIUM

TABLET, DELAYED RELEASE; ORAL

DIASONE SODIUM

ABBVIE	165MG	N006044 003	
--------	-------	-------------	--

SULFUR

POWDER; TOPICAL

BENSULFOID

POYTHRESS	33.32%	N002918 001	
-----------	--------	-------------	--

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

SULINDAC

TABLET; ORAL

CLINORIL

+ MERCK

150MG **

N017911 001

+

200MG **

N017911 002

SULINDAC

ANI PHARMS

150MG

A072973 002 Feb 28, 1992

200MG

A072973 001 Feb 28, 1992

CHARTWELL RX

150MG

A072712 001 Aug 30, 1991

200MG

A072713 001 Aug 30, 1991

EPIC PHARMA LLC

150MG

A073262 002 Sep 06, 1991

200MG

A073262 001 Sep 06, 1991

RISING

150MG

A073039 002 Jun 22, 1993

200MG

A073039 001 Jun 22, 1993

SUMATRIPTAN

SPRAY; NASAL

IMITREX

GLAXOSMITHKLINE

10MG/SPRAY

N020626 002 Aug 26, 1997

SUMATRIPTAN SUCCINATE

INJECTABLE; SUBCUTANEOUS

ALSUMA

MERIDIAN MEDCL

EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)

N022377 001 Jun 29, 2010

IMITREX

+ GLAXOSMITHKLINE

EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML) **

N020080 001 Dec 28, 1992

SUMATRIPTAN SUCCINATE

ANTARES PHARMA INC

EQ 4MG BASE/0.5ML (EQ 8MG BASE/ML)

A078319 001 Dec 10, 2015

EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)

A078319 002 Dec 10, 2015

BAXTER HLTHCARE CORP

EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)

A207101 001 Jan 19, 2023

ENDO OPERATIONS

EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)

A077332 001 Oct 09, 2009

EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)

A077871 001 Jul 09, 2009

FRESENIUS KABI USA

EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)

A079240 002 Sep 18, 2009

EQ 4MG BASE/0.5ML (EQ 8MG BASE/ML)

A079240 001 Sep 18, 2009

NORVIUM BIOSCIENCE

EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)

A203322 001 Apr 14, 2014

SANDOZ

EQ 4MG BASE/0.5ML (EQ 8MG BASE/ML)

A078067 002 Feb 06, 2009

EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)

A078067 001 Feb 06, 2009

STERISCIENCE SPECLTS

EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)

A090314 001 Jun 10, 2010

EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)

A090641 001 Jul 28, 2010

SUN PHARM

EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)

A090358 001 Jun 21, 2011

TEVA PARENTERAL

EQ 4MG BASE/0.5ML (EQ 8MG BASE/ML)

A078318 001 Feb 06, 2009

EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)

A078318 002 Feb 06, 2009

TEVA PHARMS USA

EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)

A077907 001 Feb 06, 2009

ZYDUS

EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)

A090310 001 Aug 11, 2010

SUMAVEL DOSEPRO

+ ENDO OPERATIONS

EQ 4MG BASE/0.5ML (EQ 8MG BASE/ML)

N022239 002 Nov 26, 2013

+

EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)

N022239 001 Jul 15, 2009

SYSTEM; IONTOPHORESIS

ZECURITY

+ TEVA BRANDED PHARM

EQ 6.5MG BASE/4HR

N202278 001 Jan 17, 2013

TABLET; ORAL

SUMATRIPTAN SUCCINATE

FOSUN PHARMA

EQ 25MG BASE

A076976 001 Aug 10, 2009

EQ 50MG BASE

A076976 002 Aug 10, 2009

EQ 100MG BASE

A076976 003 Aug 10, 2009

HIKMA PHARMS

EQ 25MG BASE

A078298 001 May 21, 2013

EQ 50MG BASE

A078298 002 May 21, 2013

EQ 100MG BASE

A078298 003 May 21, 2013

MYLAN

EQ 25MG BASE

A077163 001 Nov 02, 2009

EQ 50MG BASE

A077163 002 Nov 02, 2009

EQ 100MG BASE

A077163 003 Nov 02, 2009

ROXANE

EQ 25MG BASE

A078241 001 Aug 10, 2009

EQ 50MG BASE

A078241 002 Aug 10, 2009

EQ 100MG BASE

A078241 003 Aug 10, 2009

SUN PHARM INDS LTD

EQ 25MG BASE

A076554 001 Aug 10, 2009

EQ 50MG BASE

A076554 002 Aug 10, 2009

EQ 100MG BASE

A076572 001 Feb 09, 2009

TEVA

EQ 25MG BASE

A076840 001 Feb 09, 2009

EQ 50MG BASE

A076840 002 Feb 09, 2009

EQ 100MG BASE

A076840 003 Feb 09, 2009

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

SUPROFEN

SOLUTION/DROPS;OPHTHALMIC

PROFENAL

ALCON

1%

N019387 001 Dec 23, 1988

TACRINE HYDROCHLORIDE

CAPSULE;ORAL

COGNEX

SHIONOGI INC

EQ 10MG BASE

N020070 001 Sep 09, 1993

EQ 20MG BASE

N020070 002 Sep 09, 1993

EQ 30MG BASE

N020070 003 Sep 09, 1993

EQ 40MG BASE

N020070 004 Sep 09, 1993

TACROLIMUS

CAPSULE;ORAL

TACROLIMUS

HERITAGE PHARMA AVET

EQ 5MG BASE

A090402 001 Jul 01, 2010

INJECTABLE;INJECTION

TACROLIMUS

HOSPIRA

EQ 5MG BASE/ML

A203900 001 Aug 25, 2017

TADALAFIL

TABLET;ORAL

CIALIS

+ LILLY

2.5MG

N021368 004 Jan 07, 2008

TADALAFIL

CHARTWELL RX

2.5MG

A210716 001 Dec 29, 2020

5MG

A210716 002 Dec 29, 2020

10MG

A210716 003 Dec 29, 2020

20MG

A210716 004 Dec 29, 2020

NORVIUM BIOSCIENCE

5MG

A206957 001 Apr 29, 2019

20MG

A200630 001 Aug 03, 2018

RISING

2.5MG

A206956 001 Apr 29, 2019

10MG

A206956 002 Apr 29, 2019

20MG

A206956 003 Apr 29, 2019

ZYDUS PHARMS

20MG

A212515 001 Jun 06, 2023

TABLET, CHEWABLE;ORAL

CHEWTADZY

+ B BETTER

5MG

N218527 001 Jun 28, 2024

+

10MG

N218527 002 Jun 28, 2024

+

20MG

N218527 003 Jun 28, 2024

TALBUTAL

TABLET;ORAL

LOTUSATE

SANOFI AVENTIS US

120MG

N009410 005

TAMOXIFEN CITRATE

TABLET;ORAL

NOLVADEX

+ ASTRAZENECA

EQ 10MG BASE **

N017970 001

+

EQ 20MG BASE **

N017970 002 Mar 21, 1994

TAMOXIFEN CITRATE

ACTAVIS LABS FL INC

EQ 10MG BASE

A076179 001 Feb 20, 2003

EQ 20MG BASE

A076179 002 Feb 20, 2003

AEGIS PHARMS

EQ 10MG BASE

A076398 001 Mar 31, 2003

EQ 20MG BASE

A076398 002 Mar 31, 2003

IVAX SUB TEVA PHARMS

EQ 10MG BASE

A075740 001 Feb 20, 2003

EQ 20MG BASE

A075740 002 Feb 20, 2003

PHARMACHEMIE

EQ 10MG BASE

A074539 001 Mar 31, 2003

ROXANE

EQ 10MG BASE

A076027 001 Feb 20, 2003

EQ 20MG BASE

A076027 002 Feb 20, 2003

TEVA

EQ 10MG BASE

A074504 001 Apr 28, 2003

EQ 20MG BASE

A074504 002 Apr 28, 2003

TAMSULOSIN HYDROCHLORIDE

CAPSULE;ORAL

TAMSULOSIN HYDROCHLORIDE

ASCENT PHARMS INC

0.4MG

A214730 001 May 04, 2022

ENDO OPERATIONS

0.4MG

A202010 001 Jan 04, 2013

NORVIUM BIOSCIENCE

0.4MG

A090408 001 Apr 27, 2010

SUN PHARM INDS LTD

0.4MG

A090931 001 Jul 15, 2010

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

TAPENTADOL HYDROCHLORIDE

SOLUTION;ORAL

NUCYNTA

+ COLLEGIUM PHARM INC EQ 20MG BASE/ML **

N203794 001 Oct 15, 2012

TAVABOROLE

SOLUTION;TOPICAL

KERYDIN

+ ANACOR PHARMS INC 5% **

N204427 001 Jul 07, 2014

TAZAROTENE

CREAM;TOPICAL

TAZAROTENE

FOUGERA PHARMS INC 0.1%

A211175 001 Jan 28, 2019

TECHNETIUM TC-99M APCITIDE

INJECTABLE;INJECTION

ACUTECT

CIS BIO INTL SA N/A

N020887 001 Sep 14, 1998

TECHNETIUM TC-99M DEPREOTIDE

INJECTABLE;INJECTION

NEO TECT KIT

CIS BIO INTL SA N/A **

N021012 001 Aug 03, 1999

TECHNETIUM TC-99M DISOFENIN KIT

INJECTABLE;INJECTION

HEPATOLITE

SUN PHARM INDS INC N/A

N018467 001 Mar 16, 1982

TECHNETIUM TC-99M ETIDRONATE KIT

INJECTABLE;INJECTION

CINTICHEM TECHNETIUM 99M HEDSPA

GE HEALTHCARE N/A

N017653 001

MPI STANNOUS DIPHOSPHONATE

GE HEALTHCARE N/A

N017667 001

OSTEOSCAN

MALLINCKRODT N/A

N017454 001

TECHNETIUM TC 99M DIPHOSPHONATE-TIN KIT

GE HEALTHCARE N/A

N017562 001

TECHNETIUM TC-99M FERSENTETATE KIT

INJECTABLE;INJECTION

RENOTEC

BRACCO N/A

N017045 001

TECHNETIUM TC-99M GLUCEPTATE KIT

INJECTABLE;INJECTION

GLUCOSCAN

BRISTOL MYERS SQUIBB N/A

N017907 001

TECHNESCAN GLUCEPTATE

DRAXIMAGE N/A

N018272 001 Jan 27, 1982

TECHNETIUM TC-99M LIDOFENIN KIT

INJECTABLE;INJECTION

TECHNESCAN HIDA

DRAXIMAGE N/A

N018489 001 Oct 31, 1986

TECHNETIUM TC-99M MEDRONATE

INJECTABLE;INJECTION

DRAXIMAGE MDP-10

JUBILANT N/A

N018035 001

TECHNETIUM TC-99M MEDRONATE KIT

INJECTABLE;INJECTION

AMERSCAN MDP KIT

GE HEALTHCARE N/A

N018335 001 Aug 05, 1982

OSTEOLITE

PHARMALUCENCE N/A

N017972 001

TECHNETIUM TC 99M MPI MDP

GE HEALTHCARE N/A

N018141 001

N/A

N018141 002 Jun 12, 1989

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

TECHNETIUM TC-99M PENTETATE KIT

INJECTABLE; INJECTION

AN-DTPA

JUBILANT DRAXIMAGE	N/A	N017714	001
MPI DTPA KIT - CHELATE			
GE HEALTHCARE	N/A	N017255	001
TECHNETIUM TC-99M PENTETATE KIT			
GE HEALTHCARE	N/A	N017264	002

TECHNETIUM TC-99M POLYPHOSPHATE KIT

INJECTABLE; INJECTION

SODIUM POLYPHOSPHATE-TIN KIT

GE HEALTHCARE	N/A	N017664	001
---------------	-----	---------	-----

TECHNETIUM TC-99M PYRO/TRIMETA PHOSPHATES KIT

INJECTABLE; INJECTION

PYROLITE

PHARMALUCENCE	N/A	N017684	001
---------------	-----	---------	-----

TECHNETIUM TC-99M PYROPHOSPHATE KIT

INJECTABLE; INJECTION

PHOSPHOTEC

BRACCO	N/A	N017680	001
--------	-----	---------	-----

TECHNETIUM TC-99M RED BLOOD CELL KIT

INJECTABLE; INJECTION

RBC-SCAN

CADEMA	N/A	N020063	001	Jun 11, 1992
--------	-----	---------	-----	--------------

TECHNETIUM TC-99M SESTAMIBI KIT

INJECTABLE; INJECTION

MIRALUMA

LANTHEUS MEDCL	N/A	N019785	003	May 23, 1997
----------------	-----	---------	-----	--------------

TECHNETIUM TC-99M SODIUM PERTECHNETATE

SOLUTION; INJECTION, ORAL

SODIUM PERTECHNETATE TC 99M

+	GE HEALTHCARE	2-100mCi/ML **	N017471	001
+	MALLINCKRODT	10-60mCi/ML **	N017725	001
	PHARMALUCENCE	12mCi/ML	N017321	001
		24mCi/ML	N017321	002
		48mCi/ML	N017321	003

TECHNETIUM TC-99M SODIUM PERTECHNETATE GENERATOR

SOLUTION; INJECTION, ORAL

MINITEC

BRACCO	0.22-2.22 CI/GENERATOR	N017339	001
--------	------------------------	---------	-----

SOLUTION; INTRAVENOUS

TECHNELITE

LANTHEUS MEDCL	0.0083-2.7 CI/GENERATOR	N017771	001
----------------	-------------------------	---------	-----

ULTRA-TECHNEKOW FM

CURIUM	0.25-3 CI/GENERATOR	N017243	002
--------	---------------------	---------	-----

SOLUTION; INTRAVENOUS, INTRAVESICULAR, OPHTHALMIC

RADIOGENIX SYSTEM

+	NORTHSTAR MEDICAL	30-1153mCi/GENERATOR	N202158	001	Feb 08, 2018
---	-------------------	----------------------	---------	-----	--------------

SOLUTION; INTRAVENOUS, ORAL

TECHNETIUM TC 99M GENERATOR

+	GE HEALTHCARE	68-2703mCi/GENERATOR	N017693	002	Dec 13, 2013
		830-16600mCi/GENERATOR	N017693	001	

TECHNETIUM TC-99M SUCCIMER KIT

INJECTABLE; INJECTION

MPI DMSA KIDNEY REAGENT

+	GE HEALTHCARE	N/A **	N017944	001	May 18, 1982
---	---------------	--------	---------	-----	--------------

TECHNETIUM TC-99M SULFUR COLLOID

SOLUTION; INJECTION, ORAL

TECHNETIUM TC 99M SULFUR COLLOID

GE HEALTHCARE	4mCi/ML	N017456	001
---------------	---------	---------	-----

SOLUTION; ORAL

TECHNETIUM TC 99M SULFUR COLLOID

MALLINCKRODT	3mCi/ML	N017724	001
--------------	---------	---------	-----

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

TECHNETIUM TC-99M SULFUR COLLOID KIT

SOLUTION; INJECTION, ORAL

TECHNECOLL

MALLINCKRODT N/A

N017059 001

TECHNETIUM TC 99M TSC

GE HEALTHCARE N/A

N017784 001

TESULOID

BRACCO N/A

N016923 001

TECHNETIUM TC-99M TEBOROXIME KIT

INJECTABLE; INJECTION

CARDIOTEC

BRACCO N/A

N019928 001 Dec 19, 1990

TEGASEROD MALEATE

TABLET; ORAL

ZELNORM

+ ALFASIGMA EQ 2MG BASE

N021200 001 Jul 24, 2002

+ EQ 6MG BASE

N021200 002 Jul 24, 2002

TELAPREVIR

TABLET; ORAL

INCIVEK

VERTEX PHARMS 375MG

N201917 001 May 23, 2011

TELAVANCIN HYDROCHLORIDE

POWDER; INTRAVENOUS

VIBATIV

+ CUMBERLAND EQ 250MG BASE/VIAL

N022110 001 Sep 11, 2009

TELBIVUDINE

SOLUTION; ORAL

TYZEKA

NOVARTIS 100MG/5ML

N022154 001 Apr 28, 2009

TABLET; ORAL

TYZEKA

+ NOVARTIS 600MG

N022011 001 Oct 25, 2006

TELITHROMYCIN

TABLET; ORAL

KETEK

SANOFI AVENTIS US 300MG

N021144 002 Feb 09, 2005

400MG

N021144 001 Apr 01, 2004

TELMISARTAN

TABLET; ORAL

TELMISARTAN

HISUN PHARM HANGZHOU 20MG

A207843 001 Feb 19, 2019

40MG

A207843 002 Feb 19, 2019

80MG

A207843 003 Feb 19, 2019

JUBILANT GENERICS 20MG

A204164 001 Aug 22, 2016

40MG

A204164 002 Aug 22, 2016

80MG

A204164 003 Aug 22, 2016

TORRENT 20MG

A203171 001 Jul 07, 2014

40MG

A203171 002 Jul 07, 2014

80MG

A203171 003 Jul 07, 2014

TEMAZEPAM

CAPSULE; ORAL

TEMAZ

QUANTUM PHARMICS 15MG

A070564 001 Oct 15, 1985

30MG

A070547 001 Oct 15, 1985

TEMAZEPAM

AMNEAL PHARMS 7.5MG

A203482 001 May 23, 2016

15MG

A203482 002 May 23, 2016

22.5MG

A203482 003 May 23, 2016

30MG

A203482 004 May 23, 2016

AUROBINDO PHARMA USA 7.5MG

A070920 002 May 21, 2010

15MG

A070920 004 Jul 07, 1986

22.5MG

A070920 003 Jun 12, 2009

30MG

A070920 001 Jul 10, 1986

DURAMED PHARMS BARR 15MG

A071708 001 Sep 29, 1988

30MG

A071709 001 Sep 29, 1988

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

TEMAZEPAM

CAPSULE; ORAL

TEMAZEPAM

SUN PHARM INDUSTRIES	15MG	A071174	001	Jul 10, 1986
	22.5MG	A071175	002	Sep 14, 2009
	30MG	A071175	001	Jul 10, 1986
USL PHARMA	15MG	A070489	001	Jul 07, 1986
	30MG	A070490	001	Jul 07, 1986
WATSON LABS	15MG	A070383	001	Mar 23, 1987
	15MG	A071446	001	May 21, 1993
	30MG	A070384	001	Mar 23, 1987
	30MG	A071447	001	May 21, 1993

TEMOZOLOMIDE

CAPSULE; ORAL

TEMODAR

+ MERCK SHARP DOHME	5MG	N021029	001	Aug 11, 1999
	20MG	N021029	002	Aug 11, 1999
	100MG	N021029	003	Aug 11, 1999
	140MG	N021029	005	Oct 19, 2006
	180MG	N021029	006	Oct 19, 2006
	250MG	N021029	004	Aug 11, 1999

TEMOZOLOMIDE

ANI PHARMS	5MG	A203490	001	Jul 13, 2016
	20MG	A203490	002	Jul 13, 2016
	100MG	A203490	003	Jul 13, 2016
	140MG	A203490	004	Jul 13, 2016
	180MG	A203490	005	Jul 13, 2016
	250MG	A203490	006	Jul 13, 2016
APOTEX	5MG	A204159	001	Jul 05, 2018
	20MG	A204159	002	Jul 05, 2018
	100MG	A204159	003	Jul 05, 2018
	140MG	A204159	004	Jul 05, 2018
	180MG	A204159	005	Jul 05, 2018
	250MG	A204159	006	Jul 05, 2018
CHARTWELL MOLECULAR	5MG	A203898	001	Feb 10, 2016
	20MG	A203898	002	Feb 10, 2016
	100MG	A203898	003	Feb 10, 2016
	140MG	A203898	004	Feb 10, 2016
	180MG	A203898	005	Feb 10, 2016
	250MG	A203898	006	Feb 10, 2016
EXTROVIS	5MG	A205227	001	Jun 29, 2016
	20MG	A205227	002	Jun 29, 2016
	100MG	A205227	003	Jun 29, 2016
	140MG	A205227	004	Jun 29, 2016
	180MG	A205227	005	Jun 29, 2016
	250MG	A205227	006	Jun 29, 2016
HERITAGE	5MG	A078879	001	Mar 01, 2010
	20MG	A078879	002	Mar 01, 2010
	100MG	A078879	003	Mar 01, 2010
	140MG	A078879	005	Mar 01, 2010
	180MG	A078879	006	Mar 01, 2010
	250MG	A078879	004	Mar 01, 2010
NIVAGEN PHARMS INC	5MG	A213328	001	Nov 23, 2021
	20MG	A213328	002	Nov 23, 2021
	100MG	A213328	003	Nov 23, 2021
	140MG	A213328	004	Nov 23, 2021
	180MG	A213328	005	Nov 23, 2021
	250MG	A213328	006	Nov 23, 2021
WATSON LABS TEVA	5MG	A203959	001	Apr 18, 2017
	20MG	A203959	002	Apr 18, 2017
	100MG	A203959	003	Apr 18, 2017
	140MG	A203959	004	Apr 18, 2017
	250MG	A203959	005	Apr 18, 2017

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

TENAPANOR HYDROCHLORIDE

TABLET; ORAL

XPHOZAH

+ ARDELYX INC EQ 10MG BASE N213931 001 Oct 17, 2023

TENIPOSIDE

INJECTABLE; INJECTION

VUMON

+ HQ SPECLT PHARMA 10MG/ML N020119 001 Jul 14, 1992

TENOFOVIR ALAFENAMIDE FUMARATE

TABLET; ORAL

TENOFOVIR ALAFENAMIDE FUMARATE

APOTEX EQ 25MG BASE A213867 001 Mar 21, 2024

TENOFOVIR DISOPROXIL FUMARATE

TABLET; ORAL

TENOFOVIR DISOPROXIL FUMARATE

CHARTWELL RX 300MG A209550 001 Feb 26, 2018

NORVIUM BIOSCIENCE 150MG A206569 001 Nov 27, 2018

200MG A206569 002 Nov 27, 2018

250MG A206569 003 Nov 27, 2018

300MG A206569 004 Nov 27, 2018

TERAZOSIN HYDROCHLORIDE

CAPSULE; ORAL

HYTRIN

+ ABBOTT EQ 1MG BASE ** N020347 001 Dec 14, 1994

+ EQ 2MG BASE ** N020347 002 Dec 14, 1994

+ EQ 5MG BASE ** N020347 003 Dec 14, 1994

+ EQ 10MG BASE ** N020347 004 Dec 14, 1994

TERAZOSIN HYDROCHLORIDE

HIKMA EQ 1MG BASE A075498 001 Apr 12, 2001

EQ 2MG BASE A075498 002 Apr 12, 2001

EQ 5MG BASE A075498 003 Apr 12, 2001

EQ 10MG BASE A075498 004 Apr 12, 2001

NORVIUM BIOSCIENCE EQ 1MG BASE A075140 002 Feb 11, 2000

EQ 1MG BASE A075384 001 Dec 01, 2000

EQ 2MG BASE A075140 003 Feb 11, 2000

EQ 2MG BASE A075384 002 Dec 01, 2000

EQ 5MG BASE A075140 001 Feb 11, 2000

EQ 5MG BASE A075384 003 Dec 01, 2000

EQ 10MG BASE A075140 004 Feb 11, 2000

EQ 10MG BASE A075384 004 Dec 01, 2000

RANBAXY LABS LTD EQ 1MG BASE A076021 001 Aug 22, 2002

EQ 2MG BASE A076021 002 Aug 22, 2002

EQ 5MG BASE A076021 003 Aug 22, 2002

EQ 10MG BASE A076021 004 Aug 22, 2002

TABLET; ORAL

HYTRIN

ABBOTT EQ 1MG BASE N019057 001 Aug 07, 1987

EQ 2MG BASE N019057 002 Aug 07, 1987

EQ 5MG BASE N019057 003 Aug 07, 1987

EQ 10MG BASE N019057 004 Aug 07, 1987

TERAZOSIN HYDROCHLORIDE

CHARTWELL RX EQ 1MG BASE A074657 001 Apr 28, 2000

EQ 2MG BASE A074657 002 Apr 28, 2000

EQ 5MG BASE A074657 003 Apr 28, 2000

EQ 10MG BASE A074657 004 Apr 28, 2000

IVAX SUB TEVA PHARMS EQ 1MG BASE A074530 001 Apr 21, 2000

EQ 2MG BASE A074530 002 Apr 21, 2000

EQ 5MG BASE A074530 003 Apr 21, 2000

EQ 10MG BASE A074530 004 Apr 21, 2000

SANDOZ EQ 1MG BASE A074315 001 Dec 31, 1998

EQ 2MG BASE A074315 002 Dec 31, 1998

EQ 5MG BASE A074315 003 Dec 31, 1998

EQ 10MG BASE A074315 004 Dec 31, 1998

TEVA EQ 1MG BASE A074446 001 May 18, 2000

EQ 2MG BASE A074446 002 May 18, 2000

EQ 5MG BASE A074446 003 May 18, 2000

EQ 10MG BASE A074446 004 May 18, 2000

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

TERBINAFINE

GEL;TOPICAL

LAMISIL

KARO HLTHCARE 1% N020846 001 Apr 29, 1998

TERBINAFINE HYDROCHLORIDE

CREAM;TOPICAL

LAMISIL

NOVARTIS 1% ** N020192 001 Dec 30, 1992

GRANULE;ORAL

LAMISIL

+ NOVARTIS EQ 125MG BASE/PACKET N022071 001 Sep 28, 2007

+ EQ 187.5MG BASE/PACKET N022071 002 Sep 28, 2007

SOLUTION;TOPICAL

LAMISIL

KARO HLTHCARE 1% N020749 001 Oct 17, 1997

TABLET;ORAL

LAMISIL

+ NOVARTIS EQ 250MG BASE ** N020539 001 May 10, 1996

TERBINAFINE HYDROCHLORIDE

CIPLA EQ 250MG BASE A077137 001 Jul 02, 2007

GEDEON RICHTER USA EQ 250MG BASE A077065 001 Jul 02, 2007

HERITAGE PHARMA AVET EQ 250MG BASE A076377 001 Jul 02, 2007

MYLAN EQ 250MG BASE A077136 001 Jul 02, 2007

NATCO PHARMA EQ 250MG BASE A077195 001 Jul 02, 2007

ROXANE EQ 250MG BASE A077223 001 Jul 02, 2007

WOCKHARDT EQ 250MG BASE A078229 001 Jul 02, 2007

TERBUTALINE SULFATE

AEROSOL, METERED;INHALATION

BRETHAIRE

NOVARTIS 0.2MG/INH N018762 001 Aug 17, 1984

BRICANYL

SANOFI AVENTIS US 0.2MG/INH N018000 001 Mar 19, 1985

INJECTABLE;INJECTION

BRETHINE

+ PHARMACARE 1MG/ML ** N018571 001

BRICANYL

SANOFI AVENTIS US 1MG/ML N017466 001

TERBUTALINE SULFATE

CHARTWELL INJECTABLE 1MG/ML A076770 001 Apr 23, 2004

DR REDDYS 1MG/ML A076853 001 Jul 20, 2004

EPIC PHARMA LLC 1MG/ML A078151 001 Jan 07, 2008

TABLET;ORAL

BRICANYL

SANOFI AVENTIS US 2.5MG N017618 001

5MG N017618 002

TERCONAZOLE

CREAM;VAGINAL

TERAZOL 3

+ JANSSEN PHARMS 0.8% ** N019964 001 Feb 21, 1991

TERAZOL 7

+ JANSSEN PHARMS 0.4% ** N019579 001 Dec 31, 1987

SUPPOSITORY;VAGINAL

TERAZOL 3

+ JANSSEN PHARMS 80MG ** N019641 001 May 24, 1988

TERCONAZOLE

FOUGERA PHARMS 80MG A076850 001 Jul 12, 2006

TERIFLUNOMIDE

TABLET;ORAL

TERIFLUNOMIDE

AMNEAL PHARMS CO 7MG A209613 001 Sep 28, 2018

14MG A209613 002 Sep 28, 2018

BRECKENRIDGE 7MG A209583 001 Sep 24, 2021

14MG A209583 002 Sep 24, 2021

MYLAN 7MG A209702 001 Feb 28, 2020

14MG A209702 002 Feb 28, 2020

TORRENT 7MG A209697 001 Apr 04, 2024

14MG A209697 002 Apr 04, 2024

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

TERIFLUNOMIDE

TABLET; ORAL

TERIFLUNOMIDE

WATSON LABS TEVA

7MG

A209549 001 Jul 27, 2018

14MG

A209549 002 Jul 27, 2018

TERIPARATIDE

SOLUTION; SUBCUTANEOUS

FORTEO

LILLY

0.75MG/3ML (0.25MG/ML)

N021318 001 Nov 26, 2002

TERIPARATIDE ACETATE

INJECTABLE; INJECTION

PARATHAR

SANOFI AVENTIS US

200 UNITS/VIAL

N019498 001 Dec 23, 1987

TESTOLACTONE

INJECTABLE; INJECTION

TESLAC

BRISTOL MYERS SQUIBB

100MG/ML

N016119 001

TABLET; ORAL

TESLAC

BRISTOL MYERS SQUIBB

50MG

N016118 001

250MG

N016118 002

TESTOSTERONE

FILM, EXTENDED RELEASE; TRANSDERMAL

ANDRODERM

+ ABBVIE

2MG/24HR **

N020489 003 Oct 20, 2011

2.5MG/24HR

N020489 001 Sep 29, 1995

+

4MG/24HR **

N020489 004 Oct 20, 2011

5MG/24HR

N020489 002 May 02, 1997

TESTODERM

ALZA

4MG/24HR

N019762 001 Oct 12, 1993

6MG/24HR

N019762 002 Oct 12, 1993

TESTODERM TTS

ALZA

5MG/24HR

N020791 001 Dec 18, 1997

GEL; TRANSDERMAL

TESTOSTERONE

ANI PHARMS

25MG/2.5GM PACKET

N202763 001 Feb 14, 2012

50MG/5GM PACKET

N202763 002 Feb 14, 2012

PERRIGO ISRAEL

25MG/2.5GM PACKET

N203098 002 Jan 31, 2013

50MG/5GM PACKET

N203098 003 Jan 31, 2013

GEL, METERED; TRANSDERMAL

FORTESTA

+ ENDO OPERATIONS

10MG/0.5GM ACTUATION **

N021463 001 Dec 29, 2010

TESTOSTERONE

ALEMBIC

1.62% (20.25MG/1.25GM ACTUATION)

A213922 001 Mar 03, 2021

PERRIGO ISRAEL

12.5MG/1.25GM ACTUATION

N203098 001 Jan 31, 2013

INJECTABLE; INJECTION

TESTOSTERONE

DR REDDYS

100MG/ML

A086417 001 Jul 07, 1983

WATSON LABS

25MG/ML

A086420 001 May 10, 1983

50MG/ML

A086419 001 Aug 23, 1983

SOLUTION, METERED; TRANSDERMAL

AXIRON

+ ELI LILLY AND CO

30MG/1.5ML ACTUATION **

N022504 001 Nov 23, 2010

TESTOSTERONE

ALEMBIC

30MG/1.5ML ACTUATION

A212882 001 Jun 14, 2021

APOTEX

30MG/1.5ML ACTUATION

A209181 001 Nov 25, 2020

TABLET, EXTENDED RELEASE; BUCCAL

STRIANT

+ AUXILIUM PHARMS LLC

30MG

N021543 001 Jun 19, 2003

TESTOSTERONE CYPIONATE

INJECTABLE; INJECTION

DEPO-TESTOSTERONE

PFIZER

50MG/ML

A085635 001

TESTOSTERONE CYPIONATE

RISING

200MG/ML

A040652 001 Dec 11, 2006

SUN PHARM INDS LTD

100MG/ML

A201720 001 Jun 03, 2013

200MG/ML

A201720 002 Jun 03, 2013

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

TESTOSTERONE CYPIONATE

INJECTABLE; INJECTION

TESTOSTERONE CYPIONATE

WATSON LABS

100MG/ML

A084401 001

100MG/ML

A086029 001

200MG/ML

A084401 002

WATSON PHARMS INC

200MG/ML

A086030 001

TESTOSTERONE ENANTHATE

INJECTABLE; INJECTION

DELATESTRYL

ENDO PHARMS

200MG/ML **

N009165 001

+

200MG/ML **

N009165 003

TESTOSTERONE ENANTHATE

RISING

200MG/ML

A040647 001 Oct 05, 2009

WATSON LABS

100MG/ML

A083667 001

100MG/ML

A085599 001

200MG/ML

A083667 002

WATSON PHARMS INC

200MG/ML

A085598 001

TESTOSTERONE PROPIONATE

INJECTABLE; INJECTION

TESTOSTERONE PROPIONATE

BEL MAR

25MG/ML

A080741 001

50MG/ML

A080742 001

100MG/ML

A080743 001

ELKINS SINN

25MG/ML

A080276 001

LILLY

50MG/ML

A080254 002

WATSON LABS

25MG/ML

A080188 001

25MG/ML

A085490 001

50MG/ML

A080188 002

50MG/ML

A085490 002

100MG/ML

A080188 003

100MG/ML

A083595 003

TETRABENAZINE

TABLET; ORAL

TETRABENAZINE

AJANTA PHARMA LTD

12.5MG

A213621 001 Dec 04, 2020

25MG

A213621 002 Dec 04, 2020

HIKMA

12.5MG

A209739 001 Apr 08, 2019

25MG

A209739 002 Apr 08, 2019

SUN PHARM

12.5MG

A206129 001 Aug 17, 2015

25MG

A206129 002 Aug 17, 2015

TETRACYCLINE HYDROCHLORIDE

CAPSULE; ORAL

BRISTACYCLINE

BRISTOL

250MG

A061658 001

250MG

A061888 001

500MG

A061658 002

500MG

A061888 002

CYCLOPAR

WARNER CHILCOTT

250MG

A061725 001

250MG

A062175 001

250MG

A062332 001

500MG

A061725 002

500MG

A062332 002

PANMYCIN

PHARMACIA AND UPJOHN

250MG

A060347 001

RETET

SOLVAY

250MG

A061443 001

500MG

A061443 002

ROBITET

WYETH AYERST

250MG

A061734 001

500MG

A061734 002

SUMYCIN

APOTHECON

100MG

A060429 002

125MG

A060429 004

250MG

A060429 001

500MG

A060429 003

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

TETRACYCLINE HYDROCHLORIDE

CAPSULE; ORAL

TETRACHEL

ANGUS	250MG	A060343 001
	500MG	A060343 003

TETRACYCLINE HYDROCHLORIDE

ABBOTT	250MG	A061802 001
	500MG	A061802 002

ANI PHARMS	250MG	A061471 001
------------	-------	-------------

AUROBINDO PHARMA USA	250MG	A060783 001
----------------------	-------	-------------

	500MG	A060783 002
--	-------	-------------

ELKINS SINN	250MG	A060059 001
-------------	-------	-------------

FERRANTE	125MG	A060173 001
----------	-------	-------------

	250MG	A060173 002
--	-------	-------------

HEATHER	250MG	A061148 001
---------	-------	-------------

	500MG	A061148 002
--	-------	-------------

HIKMA	250MG	A060768 001
-------	-------	-------------

	500MG	A060768 002
--	-------	-------------

IMPAX LABS	100MG	A060469 002
------------	-------	-------------

	250MG	A060469 001
--	-------	-------------

	500MG	A060469 003
--	-------	-------------

IVAX SUB TEVA PHARMS	250MG	A060704 001
----------------------	-------	-------------

	500MG	A060704 002
--	-------	-------------

MAST MM	250MG	A062085 001
---------	-------	-------------

PUREPAC PHARM	250MG	A060290 001
---------------	-------	-------------

	500MG	A060290 002
--	-------	-------------

PVT FORM	250MG	A062686 001	Jul 24, 1986
----------	-------	-------------	--------------

	500MG	A062686 002	Jul 24, 1986
--	-------	-------------	--------------

ROXANE	500MG	A061214 002
--------	-------	-------------

SUN PHARM INDUSTRIES	250MG	A060736 001
----------------------	-------	-------------

	500MG	A060736 002
--	-------	-------------

SUPERPHARM	250MG	A062540 001	Mar 21, 1985
------------	-------	-------------	--------------

	500MG	A062540 002	Mar 21, 1985
--	-------	-------------	--------------

VALEANT PHARM INTL	250MG	A060471 001
--------------------	-------	-------------

	500MG	A060471 002
--	-------	-------------

WARNER CHILCOTT	250MG	A062300 001
-----------------	-------	-------------

	500MG	A062300 002
--	-------	-------------

WATSON LABS	250MG	A062103 001
-------------	-------	-------------

	250MG	A062343 001
--	-------	-------------

	500MG	A062103 002
--	-------	-------------

	500MG	A062343 002
--	-------	-------------

WYETH AYERST	250MG	A061685 001
--------------	-------	-------------

	500MG	A061685 002
--	-------	-------------

TETRACYN

PFI PHARMECS	250MG	A060082 003
--------------	-------	-------------

	500MG	A060082 004
--	-------	-------------

FIBER, EXTENDED RELEASE; PERIODONTAL

ACTISITE

SCHIFF AND CO	12.7MG/FIBER	N050653 001	Mar 25, 1994
---------------	--------------	-------------	--------------

FOR SOLUTION; TOPICAL

TOPICYCLINE

SHIRE	2.2MG/ML	N050493 001
-------	----------	-------------

INJECTABLE; INJECTION

ACHROMYCIN

LEDERLE	250MG/VIAL	N050273 002
---------	------------	-------------

	500MG/VIAL	N050273 003
--	------------	-------------

TETRACYN

PFIZER	250MG/VIAL	A060096 001
--------	------------	-------------

	500MG/VIAL	A060096 002
--	------------	-------------

OINTMENT; OPHTHALMIC

ACHROMYCIN

STORZ	10MG/GM	N050266 001
-------	---------	-------------

SUSPENSION; ORAL

ACHROMYCIN V

LEDERLE	125MG/5ML	N050263 002
---------	-----------	-------------

SUMYCIN

ENDO OPERATIONS	125MG/5ML	A060400 001
-----------------	-----------	-------------

TETRACYCLINE HYDROCHLORIDE

ALPHARMA US PHARMS	125MG/5ML	A060633 001
--------------------	-----------	-------------

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

TETRACYCLINE HYDROCHLORIDE

SUSPENSION;ORAL

TETRACYCLINE HYDROCHLORIDE

FERRANTE	125MG/5ML	A060174	001
PROTER	125MG/5ML	A060446	001
PUREPAC PHARM	125MG/5ML	A060291	001

TETRACYN

PFIPHARMECS	125MG/5ML	A060095	001
-------------	-----------	---------	-----

TETRAMED

IVAX SUB TEVA PHARMS	125MG/5ML	A061468	001
----------------------	-----------	---------	-----

SUSPENSION/DROPS;OPHTHALMIC

ACHROMYCIN

STORZ	1%	N050268	001
-------	----	---------	-----

TABLET;ORAL

PANMYCIN

PHARMACIA AND UPJOHN	250MG	A061705	001
	500MG	A061705	002

SUMYCIN

STRIDES PHARMA	50MG	A061147	003
	100MG	A061147	002

TETRACYCLINE PHOSPHATE COMPLEX

CAPSULE;ORAL

TETREX

BRISTOL	EQ 100MG HYDROCHLORIDE	A061653	001
	EQ 250MG HYDROCHLORIDE	A061653	002
	EQ 250MG HYDROCHLORIDE	A061889	002
	EQ 250MG HYDROCHLORIDE	N050212	002
	EQ 500MG HYDROCHLORIDE	A061653	003
	EQ 500MG HYDROCHLORIDE	A061889	001
	EQ 500MG HYDROCHLORIDE	N050212	003

THALIDOMIDE

CAPSULE;ORAL

THALIDOMIDE

NATCO	150MG	A213267	001	Apr 27, 2023
-------	-------	---------	-----	--------------

THALLOUS CHLORIDE TL-201

INJECTABLE;INJECTION

THALLOUS CHLORIDE TL 201

BRACCO	1mCi/ML	N018548	001	Dec 30, 1982
+ GE HEALTHCARE	1mCi/ML	N018110	002	Feb 27, 1996
+ LANTHEUS MEDCL	1mCi/ML **	N017806	001	
TRACE LIFE	1mCi/ML	A075569	001	Nov 21, 2001

INJECTABLE;INTRAVENOUS

THALLOUS CHLORIDE TL 201

CURIUM	2mCi/ML	A077698	001	Nov 09, 2006
+ LANTHEUS MEDCL	2mCi/ML **	N017806	002	Oct 09, 1998

THEOPHYLLINE

CAPSULE;ORAL

BRONKODYL

SANOFI AVENTIS US	100MG	A085264	001
	200MG	A085264	002

ELIXOPHYLLIN

FOREST LABS	100MG	A085545	001	Jul 31, 1984
	200MG	A083921	001	Jul 31, 1984

SOMOPHYLLIN-T

FISONS	100MG	A087155	001	Feb 25, 1985
	200MG	A087155	002	Feb 25, 1985
	250MG	A087155	003	Feb 25, 1985

THEOPHYLLINE

KV PHARM	100MG	A085263	001	
	200MG	A085263	002	
SCHERER RP	100MG	A084731	002	Nov 07, 1986
	200MG	A084731	001	Nov 07, 1986
	250MG	A084731	003	Nov 07, 1986

CAPSULE, EXTENDED RELEASE;ORAL

AEROLATE III

FLEMING PHARMS	65MG	A085075	003	Nov 24, 1986
----------------	------	---------	-----	--------------

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

THEOPHYLLINE

CAPSULE, EXTENDED RELEASE;ORAL

AEROLATE JR					
FLEMING PHARMS	130MG	A085075	002	Nov 24,	1986
AEROLATE SR					
FLEMING PHARMS	260MG	A085075	001	Nov 24,	1986
ELIXOPHYLLIN SR					
FOREST LABS	125MG	A086826	001	Jan 29,	1985
	250MG	A086826	002	Jan 29,	1985
SLO-BID					
SANOFI AVENTIS US	50MG	A088269	001	Jan 31,	1985
	75MG	A089539	001	May 10,	1989
	100MG	A087892	001	Jan 31,	1985
	125MG	A089540	001	May 10,	1989
	200MG	A087893	001	Jan 31,	1985
	300MG	A087894	001	Jan 31,	1985
SLO-PHYLLIN					
SANOFI AVENTIS US	60MG	A085206	001	May 24,	1982
+	125MG	A085203	001	May 24,	1982
	250MG	A085205	001	May 24,	1982
SOMOPHYLLIN-CRT					
GRAHAM DM	50MG	A087763	001	Feb 27,	1985
	100MG	A087194	001		
	200MG	A088382	001	Feb 27,	1985
+	250MG	A087193	001		
	300MG	A088383	001	Feb 27,	1985
THEO-DUR					
SCHERING	50MG	A088022	001	Sep 10,	1985
	75MG	A088015	001	Sep 10,	1985
	125MG	A088016	001	Sep 10,	1985
	200MG	A087995	001	Sep 10,	1985
THEOBID					
WHITBY	260MG	A085983	001	Mar 20,	1985
THEOBID JR.					
WHITBY	130MG	A087854	001	Mar 20,	1985
THEOCLEAR L.A.-130					
SCHWARZ PHARMA	130MG	A086569	001	May 27,	1982
THEOCLEAR L.A.-260					
SCHWARZ PHARMA	260MG	A086569	002	May 27,	1982
THEOPHYL-SR					
ORTHO MCNEIL PHARM	125MG	A086480	001	Feb 08,	1985
	250MG	A086471	001	Feb 08,	1985
THEOPHYLLINE					
CENT PHARMS	125MG	A088654	001	Feb 12,	1985
	250MG	A088689	001	Feb 12,	1985
HOSPIRA	100MG	A089976	001	Jan 04,	1995
	200MG	A089977	001	Jan 04,	1995
	300MG	A089932	001	Jan 04,	1995
INWOOD LABS	100MG	A040052	001	Feb 14,	1994
	125MG	A040052	002	Feb 14,	1994
	200MG	A040052	003	Feb 14,	1994
	300MG	A040052	004	Feb 14,	1994
SANDOZ	260MG	A087462	001	May 11,	1982
THEOPHYLLINE-SR					
SCHERER RP	300MG	A088255	001	Jun 12,	1986
THEOVENT					
SCHERING	125MG	A087010	001	Jan 31,	1985
	250MG	A087910	001	Jan 31,	1985
ELIXIR;ORAL					
ELIXOMIN					
CENCI	80MG/15ML	A088303	001	Jan 25,	1984
LANOPHYLLIN					
LANNETT	80MG/15ML	A084578	001		
THEOLIXIR					
PANRAY	80MG/15ML	A084559	001		
THEOPHYL-225					
ORTHO MCNEIL PHARM	112.5MG/15ML	A086485	001		
THEOPHYLLINE					
ALPHARMA US PHARMS	80MG/15ML	A089223	001	May 27,	1988

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

THEOPHYLLINE

ELIXIR; ORAL

THEOPHYLLINE

CENCI	80MG/15ML	A087679	001	Apr 15, 1982
CHARTWELL RX	80MG/15ML	A085952	001	
HALSEY	80MG/15ML	A085169	001	
PHARM ASSOC	80MG/15ML	A086720	001	
+ PRECISION DOSE	80MG/15ML	A085863	001	
ROXANE	80MG/15ML	A084739	001	
TARO	80MG/15ML	A089626	001	Oct 28, 1988
WOCKHARDT	80MG/15ML	A086748	001	

INJECTABLE; INJECTION

THEOPHYLLINE 0.04% AND DEXTROSE 5% IN PLASTIC CONTAINER

B BRAUN	40MG/100ML	N019083	001	Nov 07, 1984
+	40MG/100ML	N019826	001	Aug 14, 1992

THEOPHYLLINE 0.08% AND DEXTROSE 5% IN PLASTIC CONTAINER

B BRAUN	80MG/100ML	N019083	002	Nov 07, 1984
+	80MG/100ML	N019826	002	Aug 14, 1992

THEOPHYLLINE 0.16% AND DEXTROSE 5% IN PLASTIC CONTAINER

B BRAUN	160MG/100ML	N019083	003	Nov 07, 1984
+	160MG/100ML	N019826	003	Aug 14, 1992

THEOPHYLLINE 0.2% AND DEXTROSE 5% IN PLASTIC CONTAINER

B BRAUN	200MG/100ML	N019212	001	Nov 07, 1984
	200MG/100ML	N019826	004	Aug 14, 1992

THEOPHYLLINE 0.32% AND DEXTROSE 5% IN PLASTIC CONTAINER

+ B BRAUN	320MG/100ML	N019826	006	Aug 14, 1992
-----------	-------------	---------	-----	--------------

THEOPHYLLINE 0.4% AND DEXTROSE 5% IN PLASTIC CONTAINER

B BRAUN	4MG/ML	N019212	003	Nov 07, 1984
	400MG/100ML	N019212	002	Nov 07, 1984
	400MG/100ML	N019826	005	Aug 14, 1992

THEOPHYLLINE AND DEXTROSE 5% IN PLASTIC CONTAINER

BAXTER HLTHCARE	4MG/ML	N018649	007	Jul 26, 1982
	40MG/100ML	N018649	001	Jul 26, 1982
	80MG/100ML	N018649	002	Jul 26, 1982
	160MG/100ML	N018649	003	Jul 26, 1982
	200MG/100ML	N018649	004	Jul 26, 1982
	320MG/100ML	N018649	006	Nov 13, 1985
	400MG/100ML	N018649	005	Jul 26, 1982

THEOPHYLLINE IN DEXTROSE 5% IN PLASTIC CONTAINER

+ HOSPIRA INC	4MG/ML	N019211	007	Dec 14, 1984
+	40MG/100ML	N019211	001	Dec 14, 1984
	80MG/100ML	N019211	002	Dec 14, 1984
+	160MG/100ML	N019211	003	Dec 14, 1984
	200MG/100ML	N019211	004	Dec 14, 1984
+	320MG/100ML	N019211	006	Jan 20, 1988
	400MG/100ML	N019211	005	Dec 14, 1984

SOLUTION; ORAL

AEROLATE

FLEMING PHARMS	150MG/15ML	A089141	001	Dec 03, 1986
----------------	------------	---------	-----	--------------

THEOLAIR

3M	80MG/15ML	A086107	001	
----	-----------	---------	-----	--

THEOPHYLLINE

+ ROXANE	80MG/15ML	A087449	001	Sep 15, 1983
----------	-----------	---------	-----	--------------

SUSPENSION; ORAL

ELIXICON

FOREST LABS	100MG/5ML	A085502	001	
-------------	-----------	---------	-----	--

SYRUP; ORAL

ACCUREBRON

SANOFI AVENTIS US	150MG/15ML	A088746	001	Nov 22, 1985
-------------------	------------	---------	-----	--------------

AQUAPHYLLIN

FERNDALE LABS	80MG/15ML	A087917	001	Jan 18, 1983
---------------	-----------	---------	-----	--------------

SLO-PHYLLIN

SANOFI AVENTIS US	80MG/15ML	A085187	001	
-------------------	-----------	---------	-----	--

THEOCLEAR-80

CENT PHARMS	80MG/15ML	A087095	001	Mar 01, 1982
-------------	-----------	---------	-----	--------------

THEOPHYLLINE

ALPHARMA US PHARMS	80MG/15ML	A086001	001	
--------------------	-----------	---------	-----	--

+	150MG/15ML	A086545	001	
---	------------	---------	-----	--

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

THEOPHYLLINE

TABLET; ORAL

QUIBRON-T

MONARCH PHARMS 300MG A088656 001 Aug 22, 1985

SLO-PHYLLIN

SANOFI AVENTIS US 100MG A085202 001

200MG A085204 001

THEOCLEAR-100

CENT PHARMS 100MG A085353 002

THEOCLEAR-200

CENT PHARMS 200MG A085353 001

THEOLAIR

MEDICIS 125MG A086399 001

250MG A086399 002

THEOPHYL-225

ORTHO MCNEIL PHARM 225MG A084726 001

TABLET, CHEWABLE; ORAL

THEOPHYL

ORTHO MCNEIL PHARM 100MG A086506 001 Sep 12, 1985

TABLET, EXTENDED RELEASE; ORAL

DURAPHYL

FOREST LABS 100MG A088503 001 Apr 03, 1985

200MG A088504 001 Apr 03, 1985

300MG A088505 001 Apr 03, 1985

LABID

WARNER CHILCOTT 250MG A087225 001

QUIBRON-T/SR

MONARCH PHARMS 300MG A087563 001 Jun 21, 1983

SUSTAIRE

ROERIG 100MG A085665 001

300MG A085665 002

T-PHYL

PHARM RES ASSOC 200MG A088253 001 Aug 17, 1983

THEO-DUR

+ SCHERING 100MG ** A085328 001

+ 200MG A086998 001

+ 300MG ** A085328 002

450MG A089131 001 Jun 25, 1986

THEOLAIR-SR

3M 200MG A088369 001 Jul 16, 1987

250MG A086363 002 Jul 16, 1987

300MG A088364 001 Jul 16, 1987

500MG A089132 001 Jul 16, 1987

THEOPHYLLINE

ABLE 300MG A040548 001 Apr 30, 2004

400MG A040543 001 Apr 27, 2004

450MG A040546 001 Apr 30, 2004

600MG A040539 001 Apr 27, 2004

HERITAGE PHARMA AVET 100MG A089807 001 Apr 30, 1990

200MG A089808 001 Apr 30, 1990

INWOOD LABS 450MG A040034 001 Apr 28, 1995

RHODES PHARMS 600MG A040086 001 Apr 15, 1996

UNI-DUR

SCHERING 400MG A089822 001 Jan 04, 1995

600MG A089823 001 Jan 04, 1995

THEOPHYLLINE SODIUM GLYCINATE

ELIXIR; ORAL

SYNOPHYLATE

CENT PHARMS EQ 165MG BASE/15ML N006333 008

TABLET; ORAL

ASBRON

NOVARTIS EQ 150MG BASE A085148 001

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

THIABENDAZOLE

SUSPENSION; ORAL

MINTEZOL

MERCCK SHARP DOHME 500MG/5ML N016097 001

TABLET, CHEWABLE; ORAL

MINTEZOL

MERCCK SHARP DOHME 500MG N016096 001

THIAMINE HYDROCHLORIDE

INJECTABLE; INJECTION

BETALIN S

+ LILLY

100MG/ML A080853 001

THIAMINE HYDROCHLORIDE

ABRAXIS PHARM 100MG/ML A080509 001

BEL MAR 100MG/ML A080718 001

200MG/ML A080712 001

DELL LABS 100MG/ML A083775 001

EPIC PHARMA LLC 100MG/ML A087968 001 Oct 01, 1982

HOSPIRA 100MG/ML A040079 001 May 03, 1996

LUITPOLD 100MG/ML A080667 001

PARKE DAVIS 100MG/ML A080770 001

WATSON LABS 100MG/ML A083534 001

200MG/ML A083534 002

WYETH AYERST 100MG/ML A080553 001

THIAMYLAL SODIUM

INJECTABLE; INJECTION

SURITAL

PARKEDALE 1GM/VIAL N007600 003

5GM/VIAL N007600 005

10GM/VIAL N007600 009

THIETHYLPERAZINE MALATE

INJECTABLE; INJECTION

TORECAN

NOVARTIS 5MG/ML N012754 002

THIETHYLPERAZINE MALEATE

SUPPOSITORY; RECTAL

TORECAN

NOVARTIS 10MG N013247 001

TABLET; ORAL

TORECAN

NOVARTIS 10MG N012753 001

THIOPENTAL SODIUM

SUSPENSION; RECTAL

PENTOTHAL

ABBOTT 400MG/GM N011679 001

THIORIDAZINE

SUSPENSION; ORAL

MELLARIL-S

NOVARTIS EQ 25MG HYDROCHLORIDE/5ML ** N017923 001

EQ 100MG HYDROCHLORIDE/5ML ** N017923 002

THIORIDAZINE HYDROCHLORIDE

CONCENTRATE; ORAL

MELLARIL

NOVARTIS 30MG/ML ** N011808 012

100MG/ML ** N011808 018

THIORIDAZINE HYDROCHLORIDE

ACTAVIS MID ATLANTIC 100MG/ML A088229 001 Aug 23, 1983

ALPHARMA US PHARMS 30MG/ML A087766 001 Apr 26, 1983

ANI PHARMS 30MG/ML A089602 001 Nov 09, 1987

100MG/ML A089603 001 Nov 09, 1987

EPIC PHARMA LLC 30MG/ML A040125 001 Aug 16, 1996

100MG/ML A040126 001 Aug 16, 1996

PHARM ASSOC 30MG/ML A040187 001 Aug 28, 1997

100MG/ML A040213 001 May 29, 1998

SANDOZ 30MG/ML A088307 001 Nov 23, 1983

100MG/ML A088308 001 Nov 23, 1983

WOCKHARDT 30MG/ML A088258 001 Jul 25, 1983

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

THIORIDAZINE HYDROCHLORIDE

CONCENTRATE; ORAL

THIORIDAZINE HYDROCHLORIDE

100MG/ML

A088227 001 Jul 05, 1983

THIORIDAZINE HYDROCHLORIDE INTENSOL

ROXANE

30MG/ML

A088941 001 Dec 16, 1985

100MG/ML

A088942 001 Dec 16, 1985

TABLET; ORAL

MELLARIL

+ NOVARTIS

10MG **

N011808 003

+

15MG **

N011808 016

+

25MG **

N011808 006

+

50MG **

N011808 011

+

100MG **

N011808 009

+

150MG **

N011808 017

+

200MG **

N011808 015

THIORIDAZINE HYDROCHLORIDE

ANI PHARMS

10MG

A088270 001 Apr 14, 1983

10MG

A088493 001 May 17, 1985

15MG

A088271 001 Apr 14, 1983

25MG

A088272 001 Apr 14, 1983

50MG

A088194 001 Apr 14, 1983

100MG

A088273 001 Oct 03, 1983

100MG

A088456 001 May 17, 1985

CHARTWELL RX

10MG

A088131 001 Aug 30, 1983

15MG

A088132 001 Aug 30, 1983

25MG

A088133 001 Aug 30, 1983

50MG

A088134 001 Aug 30, 1983

100MG

A088135 001 Nov 20, 1984

150MG

A088136 001 Sep 17, 1986

200MG

A088137 001 Sep 17, 1986

HERITAGE PHARMA AVET

10MG

A088476 001 Nov 08, 1983

25MG

A088478 001 Nov 08, 1983

50MG

A088479 001 Nov 08, 1983

100MG

A088736 001 Jul 24, 1984

MUTUAL PHARM

10MG

A088375 001 Nov 18, 1983

25MG

A087264 001 Nov 18, 1983

50MG

A088370 001 Nov 18, 1983

100MG

A088379 001 Nov 16, 1983

MYLAN

10MG

A088332 001 Jun 27, 1983

25MG

A088333 001 Jun 27, 1983

50MG

A088334 001 Jun 27, 1983

100MG

A088335 001 Nov 18, 1983

PAR PHARM

10MG

A088351 001 Dec 05, 1983

15MG

A088352 001 Dec 05, 1983

25MG

A088336 001 Dec 05, 1983

50MG

A088322 001 Dec 05, 1983

100MG

A088480 001 Dec 29, 1983

150MG

A089764 001 Feb 09, 1988

200MG

A089765 001 Feb 09, 1988

ROXANE

10MG

A088663 001 Mar 15, 1984

25MG

A088664 001 Mar 15, 1984

50MG

A088665 001 Mar 15, 1984

100MG

A089048 001 Feb 26, 1985

SUN PHARM INDUSTRIES

10MG

A089953 004 Aug 01, 1986

15MG

A088461 001 Nov 18, 1983

25MG

A089953 003 Aug 01, 1986

50MG

A089953 002 Aug 01, 1986

100MG

A089953 001 Oct 07, 1988

150MG

A088737 001 Sep 26, 1984

200MG

A088738 001 Oct 16, 1984

SUPERPHARM

10MG

A089103 001 Jul 02, 1985

25MG

A089104 001 Jul 02, 1985

50MG

A089105 001 Jul 02, 1985

WATSON LABS

10MG

A088412 001 Sep 12, 1983

10MG

A088561 001 May 11, 1984

15MG

A088345 001 Jul 28, 1983

15MG

A088562 001 May 11, 1984

25MG

A088296 001 Jul 28, 1983

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

THIORIDAZINE HYDROCHLORIDE

TABLET; ORAL

THIORIDAZINE HYDROCHLORIDE

25MG	A088755	001	Jul 24, 1984
50MG	A088323	001	Jul 28, 1983
50MG	A088563	001	May 11, 1984
100MG	A088284	001	Aug 25, 1983
100MG	A088564	001	May 11, 1984
150MG	A088410	001	Mar 05, 1984
150MG	A088869	001	Jun 28, 1985
200MG	A088381	001	Mar 14, 1984
WATSON LABS TEVA 15MG	A088477	001	Nov 08, 1983
25MG	A088567	001	May 11, 1984
200MG	A088872	001	Apr 26, 1985
WEST WARD 10MG	A088658	001	Mar 26, 1984
15MG	A088659	001	Mar 26, 1984
25MG	A088660	001	Mar 26, 1984
50MG	A088661	001	Mar 26, 1984

THIOTEPA

INJECTABLE; INJECTION

THIOPLEX

+ IMMUNEX

15MG/VIAL **

N020058 001 Dec 22, 1994

THIOTEPA

FRESENIUS KABI USA

15MG/VIAL

A075698 001 Sep 20, 2001

IMMUNEX

15MG/VIAL

N011683 001

TEVA PARENTERAL

15MG/VIAL **

A075730 001 Apr 20, 2001

30MG/VIAL **

A075730 002 Apr 20, 2001

THIOTHIXENE

CAPSULE; ORAL

NAVANE

+ PFIZER

1MG **

N016584 001

+

2MG **

N016584 002

+

5MG **

N016584 003

+

10MG **

N016584 004

+

20MG **

N016584 005

THIOTHIXENE

AM THERAP

1MG

A071884 001 Aug 12, 1987

2MG

A071885 001 Aug 12, 1987

5MG

A071886 001 Aug 12, 1987

10MG

A071887 001 Aug 12, 1987

20MG

A072200 001 Dec 17, 1987

CHARTWELL RX

1MG

A070600 001 Jun 05, 1987

2MG

A070601 001 Jun 05, 1987

5MG

A070602 001 Jun 05, 1987

10MG

A070603 001 Jun 05, 1987

EPIC PHARMA LLC

1MG

A071529 002 Jun 24, 1987

2MG

A071529 003 Jun 24, 1987

5MG

A071529 001 Jun 24, 1987

10MG

A071529 004 Jun 24, 1987

WATSON LABS

2MG

A071626 001 Jun 25, 1987

5MG

A071627 001 Jun 25, 1987

10MG

A071628 001 Jun 25, 1987

THIOTHIXENE HYDROCHLORIDE

CONCENTRATE; ORAL

NAVANE

+ PFIZER

EQ 5MG BASE/ML

N016758 001

THIOTHIXENE HYDROCHLORIDE

ALPHARMA US PHARMS

EQ 5MG BASE/ML

A070969 001 Oct 16, 1987

PACO

EQ 1MG BASE/ML

A071917 001 Sep 20, 1989

EQ 5MG BASE/ML

A071939 001 Dec 16, 1988

TEVA

EQ 5MG BASE/ML

A071184 001 Jun 22, 1987

TEVA PHARMS

EQ 5MG BASE/ML

A071554 001 Oct 16, 1987

THIOTHIXENE HYDROCHLORIDE INTENSOL

HIKMA

EQ 5MG BASE/ML

A073494 001 Jun 30, 1992

INJECTABLE; INJECTION

NAVANE

PFIZER

EQ 2MG BASE/ML

N016904 001

EQ 10MG BASE/VIAL

N016904 002

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

TIAGABINE HYDROCHLORIDE

TABLET; ORAL

GABITRIL

CEPHALON	6MG	N020646 006	Nov 29, 2005
	8MG	N020646 007	Nov 29, 2005
	10MG	N020646 008	Nov 29, 2005
	20MG	N020646 004	Sep 30, 1997
TIAGABINE HYDROCHLORIDE			
AMNEAL PHARMS CO	2MG	A208181 001	Dec 08, 2017
	4MG	A208181 002	Dec 08, 2017
	12MG	A208181 003	Dec 08, 2017
	16MG	A208181 004	Dec 08, 2017
WILSHIRE PHARMS INC	2MG	A206857 001	Oct 13, 2017
	4MG	A206857 002	Oct 13, 2017
	12MG	A206857 003	Oct 13, 2017
	16MG	A206857 004	Oct 13, 2017

TICAGRELOR

TABLET; ORAL

TICAGRELOR

AMNEAL	90MG	A208531 001	Jan 23, 2019
MYLAN	60MG	A208597 001	Jul 09, 2021
	90MG	A208597 002	Jul 09, 2021
SIGMAPHARM LABS LLC	90MG	A208596 001	Apr 07, 2020
SUNSHINE	90MG	A208508 001	Apr 06, 2020
WATSON LABS INC	60MG	A208390 001	Sep 04, 2018
	90MG	A208390 002	Sep 04, 2018

TICARCILLIN DISODIUM

INJECTABLE; INJECTION

TICAR

GLAXOSMITHKLINE	EQ 1GM BASE/VIAL	N050497 001	
	EQ 3GM BASE/VIAL	A062690 001	Dec 19, 1986
	EQ 3GM BASE/VIAL	N050497 002	
	EQ 6GM BASE/VIAL	N050497 003	
	EQ 20GM BASE/VIAL	N050497 004	
	EQ 30GM BASE/VIAL	N050497 005	Apr 04, 1984

TICLOPIDINE HYDROCHLORIDE

TABLET; ORAL

TICLID

ROCHE PALO	125MG	N019979 001	Mar 24, 1993
	250MG	N019979 002	Oct 31, 1991
TILOPIDINE HYDROCHLORIDE			
ACTAVIS ELIZABETH	250MG	A075253 001	Aug 20, 1999
CHARTWELL RX	250MG	A075318 001	Aug 20, 1999
	250MG	A075326 001	Aug 20, 1999
MYLAN	250MG	A075161 001	Sep 13, 1999
NATCO PHARMA	250MG	A075316 001	Nov 02, 1999
SUN PHARM INDS INC	250MG	A075526 001	Sep 26, 2002
TEVA	250MG	A075149 001	Aug 20, 1999
WATSON LABS	250MG	A075309 001	Apr 26, 2000

TIGECYCLINE

POWDER; INTRAVENOUS

TIGECYCLINE

ACCORD HLTHCARE INC	50MG/VIAL	N208744 001	Jan 18, 2018
XELLIA PHARMS APS	50MG/VIAL	A205722 001	Oct 18, 2019

TILUDRONATE DISODIUM

TABLET; ORAL

SKELID

+ SANOFI AVENTIS US	EQ 200MG BASE **	N020707 001	Mar 07, 1997
---------------------	------------------	-------------	--------------

TIMOLOL

SOLUTION/DROPS; OPHTHALMIC

TIMOLOL

EPIC PHARMA LLC	0.25%	A205309 001	Sep 30, 2016
	0.5%	A205309 002	Sep 30, 2016

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

TIMOLOL MALEATE

SOLUTION, GEL FORMING/DROPS;OPHTHALMIC

TIMOLOL MALEATE

AMNEAL

EQ 0.25% BASE

A216343 001 May 23, 2024

EQ 0.5% BASE

A216343 002 May 23, 2024

SOLUTION/DROPS;OPHTHALMIC

TIMOLOL MALEATE

APOTEX INC

EQ 0.25% BASE

A075411 001 Sep 08, 2000

EQ 0.5% BASE

A075412 001 Sep 08, 2000

BAUSCH AND LOMB

EQ 0.25% BASE

A074778 001 Mar 25, 1997

EQ 0.5% BASE

A074776 001 Mar 25, 1997

EPIC PHARMA LLC

EQ 0.25% BASE

A074465 001 Mar 25, 1997

EQ 0.25% BASE

A074515 001 Mar 25, 1997

EQ 0.5% BASE

A074516 001 Mar 25, 1997

FOUGERA

EQ 0.25% BASE

A074667 001 Mar 25, 1997

EQ 0.5% BASE

A074668 001 Mar 25, 1997

HIKMA

EQ 0.5% BASE

A075163 001 Sep 10, 2002

RENOVA PHARMS

EQ 0.5% BASE

A074466 001 Mar 25, 1997

TABLET;ORAL

BLOCADREN

+ MERCK

5MG **

N018017 001

+

10MG **

N018017 002

+

20MG **

N018017 004

TIMOLOL MALEATE

ANI PHARMS

5MG

A072917 001 Jul 31, 1991

10MG

A072918 001 Jul 31, 1991

20MG

A072919 001 Jul 31, 1991

CHARTWELL RX

5MG

A072550 001 Apr 13, 1989

10MG

A072551 001 Apr 13, 1989

20MG

A072552 001 Apr 13, 1989

QUANTUM PHARMICS

5MG

A072466 001 May 19, 1989

10MG

A072467 001 May 19, 1989

20MG

A072468 001 May 19, 1989

TEVA

5MG

A072648 001 Jun 16, 1993

10MG

A072649 001 Jun 16, 1993

20MG

A072650 001 Jun 16, 1993

USL PHARMA

5MG

A072001 001 Apr 11, 1989

10MG

A072002 001 Apr 11, 1989

20MG

A072003 001 Apr 11, 1989

WATSON LABS

5MG

A072269 001 Apr 11, 1989

10MG

A072270 001 Apr 11, 1989

20MG

A072271 001 Apr 11, 1989

TINIDAZOLE

TABLET;ORAL

TINIDAZOLE

HIKMA

250MG

A201172 001 Apr 30, 2012

500MG

A201172 002 Apr 30, 2012

TINZAPARIN SODIUM

INJECTABLE;INJECTION

INNOHEP

LEO PHARMA AS

20,000 IU/ML

N020484 001 Jul 14, 2000

TIOCONAZOLE

CREAM;TOPICAL

TZ-3

PFIZER

1%

N018682 001 Feb 18, 1983

TIOPRONIN

TABLET;ORAL

TIOPRONIN

ENDO OPERATIONS

100MG

A216198 001 Jun 02, 2022

TIPRANAIVIR

SOLUTION;ORAL

APTIVUS

+ BOEHRINGER INGELHEIM

100MG/ML

N022292 001 Jun 23, 2008

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

TIROFIBAN HYDROCHLORIDE

INJECTABLE; INJECTION

AGGRASTAT

MEDICURE

EQ 12.5MG BASE/50ML (EQ 0.25MG BASE/ML)

N020912 001 May 14, 1998

SOLUTION; INTRAVENOUS

AGGRASTAT

MEDICURE

EQ 25MG BASE/500ML (EQ 0.05MG BASE/ML)

N020913 001 May 14, 1998

TIZANIDINE HYDROCHLORIDE

CAPSULE; ORAL

TIZANIDINE HYDROCHLORIDE

ENDO OPERATIONS

EQ 2MG BASE

A207199 001 Mar 14, 2017

EQ 4MG BASE

A207199 002 Mar 14, 2017

EQ 6MG BASE

A207199 003 Mar 14, 2017

NORVIUM BIOSCIENCE

EQ 2MG BASE

A091502 001 Nov 09, 2012

EQ 4MG BASE

A091502 002 Nov 09, 2012

EQ 6MG BASE

A091502 003 Nov 09, 2012

TABLET; ORAL

TIZANIDINE HYDROCHLORIDE

ANI PHARMS

EQ 2MG BASE

A076283 001 Jul 12, 2002

EQ 2MG BASE

A076284 001 Jul 03, 2002

EQ 2MG BASE

A076321 001 Sep 30, 2004

EQ 2MG BASE

A076371 001 Apr 09, 2003

EQ 4MG BASE

A076283 002 Jul 12, 2002

EQ 4MG BASE

A076284 002 Jul 03, 2002

EQ 4MG BASE

A076321 002 Sep 30, 2004

EQ 4MG BASE

A076371 002 Apr 09, 2003

AUROBINDO PHARMA USA

EQ 2MG BASE

A076282 001 Dec 16, 2003

EQ 4MG BASE

A076282 002 Dec 16, 2003

CHARTWELL RX

EQ 2MG BASE

A076280 001 Nov 26, 2002

EQ 4MG BASE

A076280 002 Jun 27, 2002

ENDO OPERATIONS

EQ 2MG BASE

A207170 001 Jan 26, 2017

EQ 4MG BASE

A207170 002 Jan 26, 2017

RISING

EQ 2MG BASE

A076354 001 Mar 28, 2003

EQ 4MG BASE

A076354 002 Mar 28, 2003

ZANAFLEX

+ LEGACY PHARMA USA

EQ 2MG BASE **

N020397 002 Feb 04, 2000

TOBRAMYCIN

SOLUTION; INHALATION

TOBRAMYCIN

LUOXIN AUROVITAS

300MG/5ML

A210871 001 Jan 22, 2021

NORVIUM BIOSCIENCE

300MG/5ML

A209554 001 Oct 13, 2017

SOLUTION/DROPS; OPHTHALMIC

AKTOB

EPIC PHARMA LLC

0.3%

A064096 001 Jan 31, 1996

TOBRAMYCIN

ALCON PHARMS LTD

0.3%

A063176 001 May 25, 1994

APOTEX INC

0.3%

A065087 001 Feb 25, 2002

TOBREX

+ NOVARTIS

0.3%

N050541 001

TOBRAMYCIN SULFATE

INJECTABLE; INJECTION

NEBCIN

LILLY

EQ 10MG BASE/ML

A062008 004

EQ 10MG BASE/ML

A062707 001 Apr 29, 1987

+

EQ 10MG BASE/ML **

N050477 005

EQ 40MG BASE/ML

A062008 001

+

EQ 1.2GM BASE/VIAL **

N050519 001

TOBRAMYCIN SULFATE

APOTHECON

EQ 10MG BASE/ML

A064021 001 May 31, 1994

EQ 40MG BASE/ML

A064021 002 May 31, 1994

EQ 40MG BASE/ML

A064026 001 May 31, 1994

EPIC PHARMA LLC

EQ 40MG BASE/ML

A205179 001 Sep 16, 2014

HIKMA

EQ 10MG BASE/ML

A063113 001 Apr 26, 1991

EQ 10MG BASE/ML

A063128 001 Nov 27, 1991

EQ 40MG BASE/ML

A063118 001 Jul 29, 1991

EQ 40MG BASE/ML

A063127 001 Nov 27, 1991

HOSPIRA

EQ 10MG BASE/ML

A063080 001 Apr 30, 1991

EQ 10MG BASE/ML

A063112 001 Apr 30, 1991

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

TOBRAMYCIN SULFATE

INJECTABLE; INJECTION

TOBRAMYCIN SULFATE

	EQ 40MG BASE/ML	A063161 001	May 29, 1991
IGI LABS INC	EQ 10MG BASE/ML	A063119 001	Oct 31, 1994
	EQ 40MG BASE/ML	A063120 001	Oct 31, 1994
	EQ 40MG BASE/ML	A063121 001	Oct 31, 1994
	EQ 40MG BASE/ML	A063122 001	Oct 31, 1994
TEVA PHARMS USA	EQ 40MG BASE/ML	A063100 001	Jan 30, 1992
WATSON LABS INC	EQ 10MG BASE/ML	A062945 001	Aug 09, 1989
	EQ 40MG BASE/ML	A062945 002	Aug 09, 1989
TOBRAMYCIN SULFATE (PHARMACY BULK)			
HOSPIRA	EQ 40MG BASE/ML **	A063116 001	May 18, 1992
TOBRAMYCIN SULFATE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER			
HOSPIRA	EQ 1.2MG BASE/ML	A063081 003	Jul 31, 1990
	EQ 1.6MG BASE/ML	A063081 006	Jun 02, 1993
	EQ 80MG BASE/100ML	A063081 001	Jul 31, 1990

TOCAINIDE HYDROCHLORIDE

TABLET; ORAL

TONOCARD

ASTRAZENECA	400MG	N018257 001	Nov 09, 1984
	600MG	N018257 002	Nov 09, 1984

TOFACITINIB CITRATE

SOLUTION; ORAL

TOFACITINIB CITRATE

HIKMA	EQ 1MG BASE/ML	A216878 001	Sep 25, 2023
-------	----------------	-------------	--------------

TABLET; ORAL

TOFACITINIB CITRATE

AJANTA PHARMA LTD	EQ 10MG BASE	A212943 001	Jun 01, 2021
MICRO LABS	EQ 5MG BASE	A209738 001	Mar 13, 2023
ZYDUS PHARMS	EQ 5MG BASE	A209829 001	Mar 13, 2023

TABLET, EXTENDED RELEASE; ORAL

TOFACITINIB

AUROBINDO PHARMA LTD	EQ 11MG BASE	A218462 001	Jun 03, 2024
ZYDUS PHARMS	EQ 11MG BASE	A214264 001	Aug 19, 2021
	EQ 22MG BASE	A214264 002	Aug 19, 2021

TOLAZAMIDE

TABLET; ORAL

TOLAZAMIDE

BARR	100MG	A070162 001	Jan 14, 1986
	250MG	A070163 001	Jan 14, 1986
	500MG	A070164 001	Jan 14, 1986
CHARTWELL RX	100MG	A071633 001	Dec 09, 1987
	250MG	A070289 001	Mar 13, 1986
	500MG	A070290 001	Mar 13, 1986
COSETTE	100MG	N018894 001	Nov 02, 1984
	250MG	N018894 002	Nov 02, 1984
	500MG	N018894 003	Nov 02, 1984
DURAMED PHARMS BARR	100MG	A070165 001	Jan 10, 1986
	250MG	A070166 001	Jan 10, 1986
	500MG	A070167 001	Jan 10, 1986
INTERPHARM	250MG	A071270 001	Sep 23, 1986
	500MG	A071271 001	Sep 23, 1986
NATCO	250MG	A070259 001	Jan 02, 1986
	500MG	A070259 003	Mar 17, 1986
PAR PHARM	100MG	A070159 001	Jan 06, 1986
	250MG	A070160 001	Jan 06, 1986
	500MG	A070161 001	Jan 06, 1986
SUN PHARM INDUSTRIES	100MG	A071357 001	Jul 16, 1987
	250MG	A071358 001	Jul 16, 1987
	500MG	A071359 001	Jul 16, 1987
SUPERPHARM	250MG	A070763 001	Jun 16, 1986
	500MG	A070764 001	Jun 16, 1986
USL PHARMA	100MG	A071355 001	Jan 11, 1988
	250MG	A070168 001	Apr 02, 1986
	500MG	A070169 001	Apr 02, 1986
WATSON LABS	100MG	A070242 001	Aug 01, 1986
	100MG	A070513 001	Jan 09, 1986

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

TOLAZAMIDETABLET; ORAL
TOLAZAMIDE

250MG	A070243	001	Aug 01, 1986
250MG	A070514	001	Jan 09, 1986
500MG	A070244	001	Aug 01, 1986
500MG	A070515	001	Jan 09, 1986

TOLINASE

+ PHARMACIA AND UPJOHN	100MG	**
+	250MG	**
+	500MG	**

N015500	002
N015500	004
N015500	005

TOLAZOLINE HYDROCHLORIDE

INJECTABLE; INJECTION

PRISCOLINE

NOVARTIS

25MG/ML

N006403 005 Feb 22, 1985

TOLBUTAMIDE

TABLET; ORAL

ORINASE

PHARMACIA AND UPJOHN	250MG	**
	500MG	**

N010670	002
N010670	001

TOLBUTAMIDE

ALRA	500MG
ANI PHARMS	500MG
ASCOT	500MG
BARR	500MG
CHARTWELL RX	500MG
DAVA PHARMS INC	500MG
NORVIUM BIOSCIENCE	500MG
PARKE DAVIS	500MG
PUREPAC PHARM	500MG
SANDOZ	500MG
SUPERPHARM	500MG
VANGARD	500MG
WATSON LABS	250MG
	500MG
	500MG
	500MG

A086141	001
A087093	001
A087541	001
Mar 01, 1983	
A087121	001
A086574	001
A086926	001
A086445	001
A086047	001
A088950	001
Jun 17, 1985	
N012678	001
A088893	001
Nov 19, 1984	
A087876	001
Apr 20, 1982	
A089110	001
May 29, 1987	
A086109	001
A087318	001
A089111	001
May 29, 1987	

TOLBUTAMIDE SODIUM

INJECTABLE; INJECTION

ORINASE DIAGNOSTIC

PHARMACIA AND UPJOHN EQ 1GM BASE/VIAL

N012095 001

TOLCAPONE

TABLET; ORAL

TASMAR

BAUSCH 200MG

N020697 002 Jan 29, 1998

TOLCAPONE

ALVOGEN	100MG
DR REDDYS LABS SA	100MG
ENDO OPERATIONS	100MG

A207729	001	Jul 29, 2020
A210095	001	Aug 01, 2019
A204584	001	Mar 26, 2015

TOLMETIN SODIUM

CAPSULE; ORAL

TOLECTIN DS

+ ORTHO MCNEIL JANSSEN EQ 400MG BASE

N018084 001

TOLMETIN SODIUM

ANI PHARMS	EQ 400MG BASE
	EQ 400MG BASE
	EQ 400MG BASE
CHARTWELL RX	EQ 400MG BASE
SUN PHARM INDUSTRIES	EQ 400MG BASE
TEVA	EQ 400MG BASE

A073308	001	Jan 24, 1992
A073392	001	Jan 24, 1992
A073519	001	May 29, 1992
A073462	001	Apr 30, 1992
A073311	001	Nov 27, 1991
A073290	001	Nov 27, 1991

TABLET; ORAL

TOLECTIN

+ ORTHO MCNEIL JANSSEN EQ 200MG BASE

N017628 001

TOLECTIN 600

+ ORTHO MCNEIL JANSSEN EQ 600MG BASE

N017628 002 Mar 08, 1989

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

TOLMETIN SODIUM

TABLET;ORAL

TOLMETIN SODIUM

ANI PHARMS	EQ 600MG BASE	A073527	001	Jun 30, 1992
CHARTWELL RX	EQ 200MG BASE	A073588	001	Jul 31, 1992
	EQ 600MG BASE	A074002	001	Sep 27, 1993
COSETTE	EQ 600MG BASE	A074399	001	Mar 28, 1996
	EQ 600MG BASE	A074729	001	Feb 27, 1997
SUN PHARM INDUSTRIES	EQ 200MG BASE	A073310	001	Nov 27, 1991

TOLTERODINE TARTRATE

CAPSULE, EXTENDED RELEASE;ORAL

TOLTERODINE TARTRATE

AUROBINDO PHARMA USA	2MG	A201486	001	Oct 31, 2013
	4MG	A201486	002	Oct 31, 2013

TABLET;ORAL

TOLTERODINE TARTRATE

APOTEX CORP	1MG	A200164	001	Sep 25, 2012
	2MG	A200164	002	Sep 25, 2012
NORVIUM BIOSCIENCE	1MG	A202641	001	Nov 27, 2012
	2MG	A202641	002	Nov 27, 2012

TOLVAPTAN

TABLET;ORAL

SAMSCA

+ OTSUKA	60MG **	N022275	003	May 19, 2009
----------	---------	---------	-----	--------------

TOPIRAMATE

CAPSULE;ORAL

TOPAMAX SPRINKLE

+ JANSSEN PHARMS	50MG **	N020844	003	Oct 26, 1998
------------------	---------	---------	-----	--------------

TOPIRAMATE

BARR	15MG	A076448	001	Apr 15, 2009
	25MG	A076448	002	Apr 15, 2009
CHARTWELL RX	15MG	A079206	001	Oct 14, 2009
	25MG	A079206	002	Oct 14, 2009
WATSON LABS	15MG	A077868	001	Apr 15, 2009
	25MG	A077868	002	Apr 15, 2009

CAPSULE, EXTENDED RELEASE;ORAL

TOPIRAMATE

AJANTA PHARMA LTD	25MG	A215663	001	Aug 15, 2023
	50MG	A215663	002	Aug 15, 2023
	100MG	A215663	003	Aug 15, 2023
	200MG	A215663	004	Aug 15, 2023
ZYDUS PHARMS	200MG	A216167	001	Feb 09, 2023

TABLET;ORAL

TOPAMAX

JANSSEN PHARMS	300MG	N020505	003	Dec 24, 1996
	400MG	N020505	006	Dec 24, 1996

TOPIRAMATE

ACTAVIS TOTOWA	25MG	A078637	001	Feb 27, 2013
	50MG	A078637	002	Feb 27, 2013
	100MG	A078637	003	Feb 27, 2013
	200MG	A078637	004	Feb 27, 2013
AIPING PHARM INC	25MG	A078499	001	Jan 07, 2010
	50MG	A078499	002	Jan 07, 2010
	100MG	A078499	003	Jan 07, 2010
	200MG	A078499	004	Jan 07, 2010
BARR	25MG	A076315	001	Mar 27, 2009
	100MG	A076315	002	Mar 27, 2009
	200MG	A076315	003	Mar 27, 2009
CHARTWELL	25MG	A078410	001	Sep 11, 2013
	50MG	A078410	002	Sep 11, 2013
	100MG	A078410	003	Sep 11, 2013
	200MG	A078410	004	Sep 11, 2013
HIKMA PHARMS	25MG	A091185	001	Nov 25, 2013
	50MG	A091185	002	Nov 25, 2013
	100MG	A091185	003	Nov 25, 2013
	200MG	A091185	004	Nov 25, 2013
NATCO	25MG	A076314	001	Mar 27, 2009
	50MG	A076314	002	Mar 27, 2009

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

TOPIRAMATETABLET; ORAL
TOPIRAMATE

	100MG	A076314 003	Mar 27, 2009
	200MG	A076314 004	Mar 27, 2009
PLIVA HRVATSKA DOO	25MG	A077905 001	Mar 30, 2009
	50MG	A077905 002	Mar 30, 2009
	100MG	A077905 003	Mar 30, 2009
	200MG	A077905 004	Mar 30, 2009
ROXANE	25MG	A076306 001	Mar 27, 2009
	50MG	A076306 002	Mar 27, 2009
	100MG	A076306 003	Mar 27, 2009
	200MG	A076306 004	Mar 27, 2009
SUN PHARM INDS LTD	25MG	A076327 001	Mar 27, 2009
	100MG	A076327 002	Mar 27, 2009
	200MG	A076327 003	Mar 27, 2009
TEVA	25MG	A076317 001	Mar 27, 2009
	50MG	A076317 002	Mar 27, 2009
	100MG	A076317 003	Mar 27, 2009
	200MG	A076317 004	Mar 27, 2009
TORRENT PHARMS	25MG	A079153 001	Mar 27, 2009
	50MG	A079153 002	Mar 27, 2009
	100MG	A079153 003	Mar 27, 2009
	200MG	A079153 004	Mar 27, 2009
WATSON LABS	25MG	A077643 001	Mar 27, 2009
	50MG	A077643 002	Mar 27, 2009
	100MG	A077643 003	Mar 27, 2009
	200MG	A077643 004	Mar 27, 2009
WOCKHARDT	25MG	A090353 001	Sep 01, 2010
	50MG	A090353 002	Sep 01, 2010
	100MG	A090353 003	Sep 01, 2010
	200MG	A090353 004	Sep 01, 2010

TOPOTECAN HYDROCHLORIDE

INJECTABLE; INJECTION

TOPOTECAN HYDROCHLORIDE

FRESENIUS KABI USA	EQ 4MG BASE/VIAL	A091376 001	Nov 29, 2010
MEITHEAL	EQ 4MG BASE/VIAL	A201166 001	Aug 08, 2012
RISING	EQ 4MG BASE/VIAL	A091542 001	Aug 28, 2012
SUN PHARM INDS LTD	EQ 4MG BASE/VIAL	A202203 001	Aug 29, 2013

SOLUTION; INTRAVENOUS

TOPOTECAN

+ SANDOZ INC	EQ 1MG BASE/ML (EQ 1MG BASE/ML) **	N200199 001	Feb 25, 2011
+	EQ 3MG BASE/3ML (EQ 1MG BASE/ML) **	N200199 002	Feb 25, 2011
+	EQ 4MG BASE/4ML (EQ 1MG BASE/ML) **	N200199 003	Feb 25, 2011

TOPOTECAN HYDROCHLORIDE

DASH PHARMS	EQ 4MG BASE/4ML (EQ 1MG BASE/ML)	A206074 001	Nov 24, 2017
MEITHEAL	EQ 4MG BASE/4ML (EQ 1MG BASE/ML)	N022453 001	Dec 20, 2012

TOREMIFENE CITRATE

TABLET; ORAL

FARESTON

+ KYOWA KIRIN	EQ 60MG BASE	N020497 001	May 29, 1997
---------------	--------------	-------------	--------------

TORSEMIDE

SOLUTION; INTRAVENOUS

DEMADEX

+ ROCHE	50MG/5ML (10MG/ML) **	N020137 002	Aug 23, 1993
+	20MG/2ML (10MG/ML) **	N020137 001	Aug 23, 1993

TORSEMIDE

AM REGENT	20MG/2ML (10MG/ML)	A090656 001	Apr 21, 2010
	50MG/5ML (10MG/ML)	A090656 002	Apr 21, 2010
HIKMA	20MG/2ML (10MG/ML)	A078007 001	Jun 11, 2008
	50MG/5ML (10MG/ML)	A078007 002	Jun 11, 2008

TABLET; ORAL

DEMADEX

+ NORVIUM BIOSCIENCE	5MG **	N020136 001	Aug 23, 1993
+	10MG **	N020136 002	Aug 23, 1993
+	20MG **	N020136 003	Aug 23, 1993
+	100MG **	N020136 004	Aug 23, 1993

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

TORSEMIDE

TABLET; ORAL

SOAANZ

+	SARFE PHARMS	20MG	N213218	001	Jun 14, 2021
+		60MG	N213218	002	Jun 14, 2021

TORSEMIDE

	SUN PHARM INDS	5MG	A078478	001	Feb 26, 2008
		10MG	A078478	002	Feb 26, 2008
		20MG	A078478	003	Feb 26, 2008
		100MG	A078478	004	Feb 26, 2008
	TEVA	5MG	A076110	001	May 14, 2002
		10MG	A076110	002	May 14, 2002
		20MG	A076110	003	May 14, 2002
		100MG	A076110	004	May 14, 2002

TRAMADOL HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

CONZIP

+	CIPHER PHARMS INC	150MG	N022370	004	Aug 01, 2011
---	-------------------	-------	---------	-----	--------------

SOLUTION; ORAL

QDOLO

+	ATHENA	5MG/ML	N214044	001	Sep 01, 2020
---	--------	--------	---------	-----	--------------

TABLET; ORAL

TRAMADOL HYDROCHLORIDE

	ACCORD HLTHCARE	50MG	A202390	001	May 16, 2013
	ACTAVIS ELIZABETH	50MG	A075960	001	Jun 19, 2002
	ASTA	50MG	A075974	001	Jul 12, 2002
	GRAVITI PHARMS	50MG	A075968	001	Jun 25, 2002
	IPCA LABS LTD	50MG	A201973	001	Nov 16, 2012
	IVAX SUB TEVA PHARMS	50MG	A075963	001	Jul 03, 2002
	MACLEODS PHARMS LTD	50MG	A205702	001	Sep 25, 2015
	MYLAN	50MG	A075986	001	Jun 21, 2002
	MYLAN PHARMS INC	50MG	A075980	001	Nov 21, 2002
	NORTHSTAR HLTHCARE	50MG	A078935	001	May 26, 2010
	PLIVA	50MG	A075982	001	Jul 01, 2002
	SPECGX LLC	50MG	A075983	001	Jun 25, 2002
	SUN PHARM INDUSTRIES	50MG	A076100	001	Jun 20, 2002
	WATSON LABS	50MG	A075962	001	Jun 24, 2002

ULTRAM

+	JANSSEN PHARMS	50MG **	N020281	002	Mar 03, 1995
+		100MG **	N020281	001	Mar 03, 1995

TABLET, EXTENDED RELEASE; ORAL

RYZOLT

+	PURDUE PHARMA	100MG **	N021745	001	Dec 30, 2008
+		200MG **	N021745	002	Dec 30, 2008
+		300MG **	N021745	003	Dec 30, 2008

TRAMADOL HYDROCHLORIDE

	ACTAVIS ELIZABETH	100MG	A091609	001	Jun 27, 2012
		200MG	A091609	002	Jun 27, 2012
		300MG	A091609	003	Jun 27, 2012
	AUROBINDO PHARMA LTD	100MG	A204421	001	Oct 20, 2015
		200MG	A204421	002	Oct 20, 2015
		300MG	A204421	003	Oct 20, 2015
	ENDO OPERATIONS	100MG	A200491	001	Jun 27, 2012
		200MG	A200491	002	Jun 27, 2012
		300MG	A200491	003	Jun 27, 2012
	MYLAN	100MG	A205257	001	Dec 22, 2015
		200MG	A205257	002	Dec 22, 2015
		300MG	A205257	003	Dec 22, 2015
	STRIDES PHARMA	100MG	A078783	001	Nov 13, 2009
		200MG	A078783	002	Nov 13, 2009
		300MG	A078783	003	Sep 20, 2011
	SUN PHARM	100MG	A091607	001	Dec 30, 2011
		100MG	A201384	001	Dec 07, 2011
		200MG	A091607	002	Dec 30, 2011
		200MG	A201384	002	Dec 07, 2011
		300MG	A091607	003	Dec 30, 2011
		300MG	A201384	003	Dec 07, 2011

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

TRAMADOL HYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL

ULTRAM ER

+	VALEANT PHARMS	100MG **	N021692 001	Sep 08, 2005
+		200MG **	N021692 002	Sep 08, 2005
+		300MG **	N021692 003	Sep 08, 2005

TABLET, ORALLY DISINTEGRATING;ORAL

RYBIX ODT

	SHIONOGI INC	50MG	N021693 001	May 05, 2005
--	--------------	------	-------------	--------------

TRAMETINIB DIMETHYL SULFOXIDE

TABLET;ORAL

MEKINIST

+	NOVARTIS	EQ 1MG **	N204114 002	May 29, 2013
---	----------	-----------	-------------	--------------

TRAMETINIB DIMETHYL SULFOXIDE

	NOVUGEN	EQ 0.5MG	A219002 001	Aug 06, 2024
		EQ 2MG	A219002 002	Aug 06, 2024

TRANDOLAPRIL

TABLET;ORAL

MAVIK

+	ABBVIE	1MG **	N020528 001	Apr 26, 1996
+		2MG **	N020528 002	Apr 26, 1996
+		4MG **	N020528 003	Apr 26, 1996

TRANDOLAPRIL

	CHARTWELL MOLECULAR	1MG	A077307 002	Jun 12, 2007
		2MG	A077307 001	Jun 12, 2007
		4MG	A077307 003	Jun 12, 2007
	DR REDDYS LABS LTD	1MG	A078493 001	Aug 25, 2008
		2MG	A078493 002	Aug 25, 2008
		4MG	A078493 003	Aug 25, 2008
	EPIC PHARMA LLC	1MG	A077256 001	Jun 12, 2007
		2MG	A077256 002	Jun 12, 2007
		4MG	A077256 003	Jun 12, 2007
	INVAGEN PHARMS	1MG	A078320 001	Jun 12, 2007
		2MG	A078320 002	Jun 12, 2007
		4MG	A078320 003	Jun 12, 2007
	NORVIUM BIOSCIENCE	1MG	A078346 001	Apr 28, 2008
		2MG	A078346 002	Apr 28, 2008
		4MG	A078346 003	Apr 28, 2008
	TEVA PHARMS	1MG	A077489 001	Dec 12, 2006
		2MG	A077489 002	Dec 12, 2006
		4MG	A077489 003	Dec 12, 2006
	WATSON LABS	1MG	A077805 001	Jun 12, 2007
		2MG	A077805 002	Jun 12, 2007
		4MG	A077805 003	Jun 12, 2007

TRANDOLAPRIL; VERAPAMIL HYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL

TARKA

+	ABBVIE	1MG;240MG **	N020591 003	Oct 22, 1996
+		2MG;180MG	N020591 001	Oct 22, 1996
+		2MG;240MG	N020591 004	Oct 22, 1996
+		4MG;240MG	N020591 002	Oct 22, 1996

TRANEXAMIC ACID

INJECTABLE;INJECTION

TRANEXAMIC ACID

	CHARTWELL RX	100MG/ML	A202755 001	Feb 25, 2016
	EPIC PHARMA LLC	100MG/ML	A206594 001	Sep 28, 2017
	RISING	100MG/ML	A206634 001	Jun 09, 2016
	ZYDUS PHARMS	100MG/ML	A205228 001	Jul 17, 2017

TABLET;ORAL

CYKLOKAPRON

	PHARMACIA AND UPJOHN	500MG	N019280 001	Dec 30, 1986
--	----------------------	-------	-------------	--------------

TRANEXAMIC ACID

	APOTEX	650MG	A202286 001	Jan 27, 2014
--	--------	-------	-------------	--------------

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

TRAVOPROST

SOLUTION/DROPS;OPHTHALMIC

IZBA

+ NOVARTIS 0.003% ** N204822 001 May 15, 2014

TRAVATAN

+ ALCON PHARMS LTD 0.004% ** N021257 001 Mar 16, 2001

TRAVOPROST

LUPIN LTD 0.004% A207040 001 May 03, 2024

TRAZODONE HYDROCHLORIDE

TABLET;ORAL

DESYREL

+ PRAGMA 50MG ** N018207 001

+ 100MG ** N018207 002

+ 150MG ** N018207 003 Mar 25, 1985

+ 300MG ** N018207 004 Nov 07, 1988

TRAZODONE HYDROCHLORIDE

AIPING PHARM INC 100MG A072483 001 Apr 30, 1990

ALVOGEN 50MG A071636 001 Apr 18, 1988

100MG A071514 001 Apr 18, 1988

AM THERAP 50MG A071139 001 Oct 29, 1986

100MG A071140 001 Oct 29, 1986

NATCO 50MG A071405 001 Feb 27, 1991

100MG A071406 001 Feb 27, 1991

NORVIUM BIOSCIENCE 50MG A090514 001 Jun 02, 2009

100MG A090514 002 Jun 02, 2009

150MG A090514 003 Jun 02, 2009

300MG A090514 004 Jun 02, 2009

QUANTUM PHARMICS 100MG A070921 001 Dec 01, 1986

RISING 50MG A072484 001 Apr 30, 1990

TEVA 150MG A074357 001 Apr 30, 1997

USL PHARMA 50MG A070491 001 Apr 29, 1987

100MG A070492 001 Apr 29, 1987

WATSON LABS 50MG A070857 001 Oct 10, 1986

50MG A071112 001 Nov 17, 1986

100MG A070858 001 Oct 10, 1986

100MG A071113 001 Nov 17, 1986

TRIALODINE

QUANTUM PHARMICS 50MG A070942 001 Dec 01, 1986

TABLET, EXTENDED RELEASE;ORAL

OLEPTRO

+ ANGELINI PHARMA 150MG ** N022411 001 Feb 02, 2010

+ 300MG ** N022411 002 Feb 02, 2010

TREPROSTINIL

SOLUTION;INTRAVENOUS, SUBCUTANEOUS

REMODULIN

UNITED THERAP 20MG/20ML (1MG/ML) N208276 001 Jul 30, 2018

50MG/20ML (2.5MG/ML) N208276 002 Jul 30, 2018

100MG/20ML (5MG/ML) N208276 003 Jul 30, 2018

200MG/20ML (10MG/ML) N208276 004 Jul 30, 2018

TRETINOIN

CAPSULE;ORAL

VESANOID

+ CHEPLAPHARM 10MG ** N020438 001 Nov 22, 1995

CREAM;TOPICAL

RENOVA

+ VALEANT PHARMS NORTH 0.05% ** N019963 001 Dec 29, 1995

TRETINOIN

ALLERGAN 0.0375% A090098 001 Mar 22, 2010

0.075% A202209 001 Oct 11, 2012

ZO SKIN HEALTH 0.05% A076498 001 Sep 15, 2005

GEL;TOPICAL

AVITA

NORVIUM BIOSCIENCE 0.025% N020400 001 Jan 29, 1998

TRETINOIN

NORVIUM BIOSCIENCE 0.04% A202567 001 Jul 17, 2013

0.1% A202026 001 Jul 17, 2013

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

TRETINOIN

SOLUTION; TOPICAL

RETIN-A

+ VALEANT INTL

0.05%

N016921 001

TRETINOIN

TEVA PHARMS

0.05%

A074873 001 Jun 19, 1998

WOCKHARDT

0.05%

A075260 001 Jan 25, 1999

SWAB; TOPICAL

RETIN-A

VALEANT INTL

0.05%

N016921 002

TRIAMCINOLONE

TABLET; ORAL

ARISTOCORT

ASTELLAS

1MG

N011161 009

2MG

N011161 004

4MG

N011161 007

8MG

N011161 011

16MG

N011161 010

KENACORT

DELCOR ASSET CORP

1MG

N011283 003

2MG

N011283 008

4MG

N011283 006

8MG

N011283 010

TRIAMCINOLONE

BARR

2MG

A084286 001

2MG

A084318 001

4MG

A084267 001

4MG

A084319 001

8MG

A084268 001

8MG

A084320 001

4MG

A084340 001

IMPAX LABS

4MG

A083750 001

IVAX SUB TEVA PHARMS

4MG

A084406 001

MYLAN

2MG

A084406 001

PUREPAC PHARM

2MG

A084020 002

4MG

A084020 003

ROXANE

2MG

A084708 001

4MG

A084709 001

8MG

A084707 001

SANDOZ

4MG

A085601 001

TEVA

4MG

A084775 001

WATSON LABS

4MG

A084270 001

4MG

A085834 001

TRIAMCINOLONE ACETONIDE

AEROSOL, METERED; INHALATION

AZMACORT

ABBVIE

0.1MG/INH

N018117 001 Apr 23, 1982

AEROSOL, METERED; NASAL

NASACORT

CHATTEM SANOFI

0.055MG/INH

N019798 001 Jul 11, 1991

CREAM; TOPICAL

ARISTOCORT

ASTELLAS

0.025%

A083017 003

0.1%

A083016 004

+

0.5%

A083015 002

ARISTOCORT A

ASTELLAS

0.025%

A083017 004

0.025%

A088818 001 Oct 16, 1984

0.1%

A083016 005

0.1%

A088819 001 Oct 16, 1984

0.5%

A083015 003

0.5%

A088820 001 Oct 16, 1984

FLUTEX

IVAX PHARMS

0.025%

A085539 001

0.1%

A085539 002

0.5%

A085539 003

KENALOG

+ DELCOR ASSET CORP

0.5%

A083943 001

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

TRIAMCINOLONE ACETONIDE

CREAM;TOPICAL

KENALOG-H			
	DELCOR ASSET CORP	0.1%	A086240 001
TRIACET			
	TEVA	0.025%	A084908 001
		0.1%	A084908 002
		0.5%	A084908 003
TRIACORT			
	SOLVAY	0.1%	A087113 001
TRIAMCINOLONE ACETONIDE			
	ACTAVIS MID ATLANTIC	0.1%	A087798 001 Jun 04, 1982
	ALPHARMA US PHARMS	0.025%	A087797 001 Jun 07, 1982
	AMBIX	0.025%	A087932 001 May 09, 1983
	MORTON GROVE	0.025%	A088094 001 Sep 01, 1983
		0.1%	A088095 001 Sep 01, 1983
		0.5%	A088096 001 Sep 01, 1983
+	NORVIUM BIOSCIENCE	0.025% **	N011601 003
+		0.1% **	N011601 006
	PHARMADERM	0.025%	A087990 001 Jul 07, 1983
		0.1%	A087991 001 Jul 07, 1983
		0.5%	A087992 001 Jul 07, 1983
	PHARMAFAIR	0.025%	A087921 001 Aug 10, 1982
		0.1%	A087912 001 Aug 10, 1982
		0.5%	A087922 001 Aug 10, 1982
	TARO	0.025%	A040038 001 Oct 26, 1994
		0.025%	A086277 001
		0.1%	A086276 001
		0.5%	A086275 001
	TOPIDERM	0.025%	A089274 001 Feb 21, 1989
		0.1%	A089275 001 Feb 21, 1989
		0.5%	A089276 001 Feb 21, 1989
TRIALEX			
	IVAX PHARMS	0.025%	A087430 001 Nov 01, 1988
		0.1%	A087429 001 Nov 01, 1988
		0.5%	A087428 001 Nov 01, 1988
TRYMEX			
	SAVAGE LABS	0.025%	A088196 001 Mar 25, 1983
		0.1%	A088197 001 Mar 25, 1983
		0.5%	A088198 001 Mar 25, 1983
GEL;TOPICAL			
ARISTOGEL			
	ASTELLAS	0.1%	A083380 001
INJECTABLE;INJECTION			
TRIAMCINOLONE ACETONIDE			
	PARNELL	3MG/ML	N019503 001 Oct 16, 1987
	SANDOZ	10MG/ML	A090166 001 May 27, 2009
		40MG/ML	A090164 001 Jun 01, 2009
	WATSON LABS	40MG/ML	A085825 001
INJECTABLE;INTRA-ARTICULAR, INTRAMUSCULAR, INTRAVITREAL			
TRIVARIS			
+	ALLERGAN	8MG/0.1ML (8MG/0.1ML) **	N022220 001 Jun 16, 2008
LOTION;TOPICAL			
KENALOG			
	DELCOR ASSET CORP	0.025% **	A084343 001
+		0.025% **	N011602 003
		0.1% **	A084343 002
+		0.1% **	N011602 001
TRIAMCINOLONE ACETONIDE			
	ALPHARMA US PHARMS	0.025%	A087191 001 Sep 08, 1982
		0.1%	A087192 001 Sep 08, 1982
	PAI HOLDINGS PHARM	0.025%	A204608 001 Jul 07, 2016
		0.1%	A204606 001 Jul 07, 2016
OINTMENT;TOPICAL			
ARISTOCORT			
	ASTELLAS	0.1%	A080750 004
+		0.5% **	A080745 002
ARISTOCORT A			
	ASTELLAS	0.1%	A080750 003

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

TRIAMCINOLONE ACETONIDE

OINTMENT; TOPICAL

ARISTOCORT A				
	0.1%		A088780 001	Oct 01, 1984
+	0.5% **		A080745 003	
	0.5%		A088781 001	Oct 05, 1984
FLUTEX				
IVAX PHARMS	0.025%		A087375 001	Nov 01, 1988
	0.1%		A087377 001	Nov 01, 1988
	0.5%		A087376 001	Nov 01, 1988
KENALOG				
+ DELCOR ASSET CORP	0.5% **		A083944 001	
TRIAMCINOLONE ACETONIDE				
ACTAVIS MID ATLANTIC	0.1%		A087799 001	Jun 07, 1982
ALPHARMA US PHARMS	0.5%		A089913 001	Dec 23, 1988
AUROBINDO PHARMA LTD	0.1%		A211315 001	Mar 18, 2020
	0.5%		A211315 002	Mar 18, 2020
COSETTE	0.5%		A208925 001	Oct 06, 2017
+ EXTROVIS	0.025% **		N011600 003	
+ MORTON GROVE	0.1% **		N011600 001	
	0.025%		A088090 001	Sep 01, 1983
	0.1%		A088091 001	Sep 01, 1983
	0.5%		A088092 001	Sep 01, 1983
PAI HOLDINGS PHARM	0.5%		A208590 001	Mar 03, 2017
PHARMADERM	0.025%		A088692 001	Aug 02, 1984
	0.1%		A088690 001	Aug 02, 1984
TARO	0.025%		A040040 001	Sep 30, 1994
	0.025%		A040374 001	Jun 05, 2001
	0.1%		A087902 001	Dec 27, 1982
	0.5%		A040386 001	Jun 05, 2001
TRIANEX				
CMP PHARMA INC	0.05%		A089595 001	Mar 23, 1995
TRYMEX				
SAVAGE LABS	0.025%		A088693 001	Aug 02, 1984
	0.1%		A088691 001	Aug 02, 1984
PASTE; DENTAL				
KENALOG IN ORABASE				
+ DELCOR ASSET CORP	0.1% **		N012097 001	
ORALONE				
TARO	0.1%		A071383 001	Jul 06, 1987
TRIAMCINOLONE ACETONIDE				
SCIEGEN PHARMS INC	0.1%		A206312 001	Aug 11, 2016
SPRAY; TOPICAL				
KENALOG				
+ SUN PHARM INDS INC	0.147MG/GM **		N012104 001	
SPRAY, METERED; NASAL				
ALLERNAZE				
LUPIN ATLANTIS	0.05MG/SPRAY		N020120 001	Feb 04, 2000
NASACORT HFA				
SANOFI AVENTIS US	0.055MG/SPRAY		N020784 001	Apr 07, 2004
TRIAMCINOLONE ACETONIDE				
PERRIGO PHARMA INTL	0.055MG/SPRAY		A078104 001	Jul 30, 2009

TRIAMCINOLONE DIACETATE

INJECTABLE; INJECTION

ARISTOCORT				
FOSUN PHARMA	25MG/ML		N011685 003	
+ TRIAMCINOLONE DIACETATE	40MG/ML **		N012802 001	
EPIC PHARMA LLC	25MG/ML		A085122 001	
	40MG/ML		A086394 001	
WATSON LABS	40MG/ML		A084072 001	
	40MG/ML		A085529 001	
SYRUP; ORAL				
ARISTOCORT				
ASTELLAS	2MG/5ML		N011960 004	
KENACORT				
DELCOR ASSET CORP	EQ 4MG BASE/5ML		N012515 001	

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

TRIAMCINOLONE HEXACETONIDE

INJECTABLE; INJECTION

ARISTOSPAN

+	ANIKA THERAPS INC	5MG/ML **	N016466	001	
+		20MG/ML **	N016466	002	

TRIAZOLAM

TABLET; ORAL

HALCION

	PFIZER	0.5MG	N017892	002	Nov 15, 1982
--	--------	-------	---------	-----	--------------

TRIAZOLAM

	HIKMA	0.125MG	A074224	001	Jun 01, 1994
		0.25MG	A074224	002	Jun 01, 1994
	NORVIUM BIOSCIENCE	0.125MG	A074031	001	Mar 25, 1994
		0.25MG	A074031	002	Mar 25, 1994
	WATSON LABS	0.125MG	A074445	001	Oct 20, 1995
		0.25MG	A074445	002	Oct 20, 1995

TRICHLORMETHIAZIDE

TABLET; ORAL

METAHYDRIN

	SANOFI AVENTIS US	2MG	N012594	001	Jun 16, 1988
		4MG	N012594	002	Jun 16, 1988

NAQUA

	SCHERING	2MG	N012265	001	
		4MG	N012265	002	

TRICHLOREX

	LANNETT	4MG	A083436	001	
		4MG	A085630	001	

TRICHLORMAS

	MAST MM	4MG	A086259	001	
--	---------	-----	---------	-----	--

TRICHLORMETHIAZIDE

	CHARTWELL RX	4MG	A085568	001	
	IMPAX LABS	4MG	A083967	001	
	PAR PHARM	2MG	A087007	001	
		4MG	A087005	001	
	SANDOZ	4MG	A086171	001	
	WATSON LABS	2MG	A083847	001	
		2MG	A086458	001	
		4MG	A083462	001	
		4MG	A083855	001	
		4MG	A085962	001	

TRICLOFOS SODIUM

SOLUTION; ORAL

TRICLOS

	SANOFI AVENTIS US	1.5GM/15ML	N016830	001	
--	-------------------	------------	---------	-----	--

TABLET; ORAL

TRICLOS

	SANOFI AVENTIS US	750MG	N016809	002	
--	-------------------	-------	---------	-----	--

TRIDIHEXETHYL CHLORIDE

INJECTABLE; INJECTION

PATHILON

	LEDERLE	10MG/ML	N009729	001	
--	---------	---------	---------	-----	--

TABLET; ORAL

PATHILON

	LEDERLE	25MG	N009489	005	
--	---------	------	---------	-----	--

TRIENTINE HYDROCHLORIDE

CAPSULE; ORAL

CLOVIQUE

	CHARTWELL RX	250MG	A209731	001	Oct 21, 2019
--	--------------	-------	---------	-----	--------------

TRIENTINE HYDROCHLORIDE

	ACCORD HLTHCARE	250MG	A212929	001	Aug 30, 2021
	AMNEAL	250MG	A210619	001	Feb 08, 2019
	CHARTWELL RX	250MG	A209415	001	Sep 16, 2019
	LUPIN LTD	250MG	A211637	001	May 21, 2020

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

TRIFLUOPERAZINE HYDROCHLORIDE

CONCENTRATE; ORAL

STELAZINE

+ GLAXOSMITHKLINE EQ 10MG BASE/ML ** N011552 006

TRIFLUOPERAZINE HYDROCHLORIDE

CHARTWELL RX EQ 10MG BASE/ML A085787 001 Apr 15, 1982

WOCKHARDT EQ 10MG BASE/ML A088143 001 Jul 26, 1983

INJECTABLE; INJECTION

STELAZINE

+ GLAXOSMITHKLINE EQ 2MG BASE/ML ** N011552 005

TABLET; ORAL

STELAZINE

+ GLAXOSMITHKLINE EQ 1MG BASE ** N011552 001

+ EQ 2MG BASE ** N011552 002

+ EQ 5MG BASE ** N011552 003

+ EQ 10MG BASE ** N011552 004

TRIFLUOPERAZINE HYDROCHLORIDE

ATHEM EQ 1MG BASE A040153 001 Oct 25, 1996

EQ 2MG BASE A040153 002 Oct 25, 1996

EQ 5MG BASE A040153 003 Oct 25, 1996

EQ 10MG BASE A040153 004 Oct 25, 1996

DURAMED PHARMS BARR EQ 1MG BASE A088967 001 Apr 23, 1985

EQ 2MG BASE A088968 001 Apr 23, 1985

EQ 5MG BASE A088969 001 Apr 23, 1985

EQ 10MG BASE A088970 001 Apr 23, 1985

IVAX PHARMS EQ 1MG BASE A087612 001 Nov 19, 1982

EQ 2MG BASE A087613 001 Nov 19, 1982

EQ 5MG BASE A087328 001 Nov 19, 1982

EQ 10MG BASE A087614 001 Nov 19, 1982

WATSON LABS EQ 1MG BASE A085975 001 Jun 23, 1988

EQ 2MG BASE A085976 001 Jun 23, 1988

EQ 5MG BASE A085973 001 Jun 23, 1988

EQ 10MG BASE A088710 001 Jun 23, 1988

TRIFLUPROMAZINE

SUSPENSION; ORAL

VESPRIN

APOTHECON EQ 50MG HYDROCHLORIDE/5ML N011491 004

TRIFLUPROMAZINE HYDROCHLORIDE

INJECTABLE; INJECTION

VESPRIN

APOTHECON 3MG/ML N011325 005

10MG/ML N011325 004

20MG/ML N011325 001

TABLET; ORAL

VESPRIN

BRISTOL MYERS SQUIBB 10MG N011123 001

25MG N011123 002

50MG N011123 003

TRIFLURIDINE

SOLUTION/DROPS; OPHTHALMIC

TRIFLURIDINE

HIKMA 1% A205438 001 Jul 28, 2017

TRIHENXYPHENIDYL HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

ARTANE

LEDERLE 5MG N006773 010

5MG N012947 001

ELIXIR; ORAL

ARTANE

LEDERLE 2MG/5ML N006773 009

TRIHENXYPHENIDYL HYDROCHLORIDE

PHARM VENTURES 2MG/5ML A089514 001 Apr 07, 1989

TABLET; ORAL

ARTANE

+ LEDERLE 2MG ** N006773 005

+ 5MG ** N006773 003

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

TRIHEXYPHENIDYL HYDROCHLORIDE

TABLET; ORAL

TREMIM

SCHERING	2MG	A080381	001	
	5MG	A080381	003	

TRIHEXYPHENIDYL HYDROCHLORIDE

HIKMA	2MG	A040337	002	Feb 16, 2000
	5MG	A040337	001	Feb 16, 2000
NYLOS	5MG	A085622	001	
VANGARD	2MG	A088035	001	Jul 30, 1982
WATSON LABS	2MG	A040184	001	Feb 06, 1998
	2MG	A085117	001	
	5MG	A040184	002	Feb 06, 1998
	5MG	A085105	001	

TRILOSTANE

CAPSULE; ORAL

MODRASTANE

BIOENVISION	30MG	N018719	002	Dec 31, 1984
	60MG	N018719	001	Dec 31, 1984

TRIMEPRAZINE TARTRATE

CAPSULE, EXTENDED RELEASE; ORAL

TEMARIL

ALLERGAN HERBERT	EQ 5MG BASE	N011316	004	
------------------	-------------	---------	-----	--

SYRUP; ORAL

TEMARIL

ALLERGAN HERBERT	EQ 2.5MG BASE/5ML	N011316	003	
------------------	-------------------	---------	-----	--

TRIMEPRAZINE TARTRATE

ALPHARMA US PHARMS	EQ 2.5MG BASE/5ML	A085015	001	Feb 18, 1982
MORTON GROVE	EQ 2.5MG BASE/5ML	A088285	001	Apr 11, 1985

TABLET; ORAL

TEMARIL

ALLERGAN HERBERT	EQ 2.5MG BASE	N011316	001	
------------------	---------------	---------	-----	--

TRIMETHADIONE

CAPSULE; ORAL

TRIDIONE

ABBVIE	300MG	N005856	005	
--------	-------	---------	-----	--

SOLUTION; ORAL

TRIDIONE

ABBVIE	200MG/5ML	N005856	002	
--------	-----------	---------	-----	--

TABLET; ORAL

TRIDIONE

+ ABBVIE	150MG	N005856	009	
----------	-------	---------	-----	--

TRIMETHAPHAN CAMSYLATE

INJECTABLE; INJECTION

ARFONAD

ROCHE	50MG/ML	N008983	001	
-------	---------	---------	-----	--

TRIMETHOBENZAMIDE HYDROCHLORIDE

CAPSULE; ORAL

TIGAN

+ KING PHARMS LLC	300MG **	N017531	006	Dec 13, 2001
-------------------	----------	---------	-----	--------------

TRIMETHOBENZAMIDE HYDROCHLORIDE

SUN PHARM INDUSTRIES	300MG	A076570	001	Aug 28, 2003
----------------------	-------	---------	-----	--------------

INJECTABLE; INJECTION

TRIMETHOBENZAMIDE HYDROCHLORIDE

AM REGENT	100MG/ML	A091330	001	Mar 08, 2011
HOSPIRA	100MG/ML	A088804	001	Apr 03, 1987
SMITH AND NEPHEW	100MG/ML	A088960	001	Apr 04, 1986
	100MG/ML	A089043	001	Apr 04, 1986
SOLOPAK	100MG/ML	A089094	001	Apr 04, 1986
WATSON LABS	100MG/ML	A086577	001	Oct 19, 1982
	100MG/ML	A087939	001	Dec 28, 1982

TRIMETHOBENZAMIDE HYDROCHLORIDE PRESERVATIVE FREE

AM REGENT	100MG/ML	A091329	001	Mar 08, 2011
-----------	----------	---------	-----	--------------

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

TRIMETHOPRIM

TABLET; ORAL

PROLOPRIM

MONARCH PHARMS	100MG	N017943 001	
	200MG	N017943 003	Jul 14, 1982

TRIMETHOPRIM

SUN PHARM INDUSTRIES	100MG	A070494 001	Jan 22, 1986
	200MG	A070495 001	Sep 24, 1986
TEVA	200MG **	A071259 001	Jun 18, 1987

TRIMPEX

ROCHE	100MG	N017952 001	
-------	-------	-------------	--

TRIMPEX 200

ROCHE	200MG	N017952 002	Nov 09, 1982
-------	-------	-------------	--------------

TRIMETHOPRIM HYDROCHLORIDE

SOLUTION; ORAL

PRIMSOL

PANGEA	EQ 25MG BASE/5ML	N074374 001	Jun 23, 1995
+	EQ 50MG BASE/5ML	N074973 001	Jan 24, 2000

TRIMETREXATE GLUCURONATE

INJECTABLE; INJECTION

NEUTREXIN

MEDIMMUNE ONCOLOGY	EQ 25MG BASE/VIAL	N020326 001	Dec 17, 1993
	EQ 200MG BASE/VIAL	N020326 002	Jul 31, 1998

TRIMIPRAMINE MALEATE

CAPSULE; ORAL

SURMONTIL

+ ODYSSEY PHARMS	EQ 25MG BASE **	N016792 001	
+	EQ 50MG BASE **	N016792 002	
+	EQ 100MG BASE **	N016792 003	Sep 15, 1982

TRIMIPRAMINE MALEATE

USL PHARMA	EQ 25MG BASE	A071283 001	Dec 08, 1987
	EQ 50MG BASE	A071284 001	Dec 08, 1987
	EQ 100MG BASE	A071285 001	Dec 08, 1987

TRIOXSALEN

TABLET; ORAL

TRISORALEN

VALEANT PHARM INTL	5MG	N012697 001	
--------------------	-----	-------------	--

TRIPLENNAMINE CITRATE

ELIXIR; ORAL

PBZ

NOVARTIS	EQ 25MG HYDROCHLORIDE/5ML	N005914 004	
----------	---------------------------	-------------	--

TRIPLENNAMINE HYDROCHLORIDE

TABLET; ORAL

PBZ

NOVARTIS	25MG	A083149 001	
	50MG	N005914 002	

TRIPLENNAMINE HYDROCHLORIDE

ANABOLIC	50MG	A083037 001	
BARR	50MG	A080744 001	
HEATHER	50MG	A083989 001	
IMPAX LABS	50MG	A080785 001	
LANNETT	50MG	A083557 001	
NYLOS	50MG	A085412 001	
PARKE DAVIS	25MG	A083625 001	
	50MG	A083626 001	
WATSON LABS	50MG	A080713 001	
	50MG	A080790 001	
	50MG	A085188 001	

TABLET, EXTENDED RELEASE; ORAL

PBZ-SR

NOVARTIS	50MG	N010533 002	
	100MG	N010533 001	

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

TRIPLE SULFA (SULFABENZAMIDE;SULFACETAMIDE;SULFATHIAZOLE)

CREAM;VAGINAL

GYNE-SULF

COSETTE	3.7%;2.86%;3.42%	A088607 001	Jun 09, 1986
---------	------------------	-------------	--------------

SULTRIN

ORTHO MCNEIL PHARM	3.7%;2.86%;3.42%	N005794 001	
--------------------	------------------	-------------	--

TRIPLE SULFA

ALPHARMA US PHARMS	3.7%;2.86%;3.42%	A087864 001	Sep 01, 1982
--------------------	------------------	-------------	--------------

FOUGERA	3.7%;2.86%;3.42%	A086424 001	
---------	------------------	-------------	--

PADAGIS US	3.7%;2.86%;3.42%	A087285 001	Nov 15, 1982
------------	------------------	-------------	--------------

TRYSUL

SAVAGE LABS	3.7%;2.86%;3.42%	A087887 001	Jul 23, 1982
-------------	------------------	-------------	--------------

VAGILIA

COSETTE	3.7%;2.86%;3.42%	A088821 001	Nov 09, 1987
---------	------------------	-------------	--------------

TABLET;VAGINAL

SULTRIN

ORTHO MCNEIL PHARM	184MG;143.75MG;172.5MG	N005794 002	
--------------------	------------------------	-------------	--

TRIPLE SULFA

FOUGERA	184MG;143.75MG;172.5MG	A088463 001	Jan 03, 1985
---------	------------------------	-------------	--------------

PHARMADERM	184MG;143.75MG;172.5MG	A088462 001	Jan 03, 1985
------------	------------------------	-------------	--------------

TRIPROLIDINE HYDROCHLORIDE

SYRUP;ORAL

ACTIDIL

GLAXOSMITHKLINE	1.25MG/5ML	N011496 002	Jul 01, 1983
-----------------	------------	-------------	--------------

MYIDYL

USL PHARMA	1.25MG/5ML	A087963 001	Jan 18, 1983
------------	------------	-------------	--------------

TRIPROLIDINE HYDROCHLORIDE

ALPHARMA US PHARMS	1.25MG/5ML	A085940 001	
--------------------	------------	-------------	--

HALSEY	1.25MG/5ML	A088735 001	Jan 17, 1985
--------	------------	-------------	--------------

PHARM ASSOC	1.25MG/5ML	A087514 001	Feb 10, 1982
-------------	------------	-------------	--------------

TABLET;ORAL

ACTIDIL

GLAXOSMITHKLINE	2.5MG	N011110 002	Jul 01, 1983
-----------------	-------	-------------	--------------

TRIPROLIDINE HYDROCHLORIDE

VITARINE	2.5MG	A085610 001	
----------	-------	-------------	--

WATSON LABS	2.5MG	A085094 001	
-------------	-------	-------------	--

TRISULFAPYRIMIDINES (SULFADIAZINE;SULFAMERAZINE;SULFAMETHAZINE)

SUSPENSION;ORAL

LANTRISUL

LANNETT	167MG/5ML;167MG/5ML;167MG/5ML	A080123 002	
---------	-------------------------------	-------------	--

NEOTRIZINE

LILLY	167MG/5ML;167MG/5ML;167MG/5ML	N006317 012	
-------	-------------------------------	-------------	--

SULFALOID

FOREST PHARMS	167MG/5ML;167MG/5ML;167MG/5ML	A080100 001	
---------------	-------------------------------	-------------	--

SULFOSE

WYETH AYERST	167MG/5ML;167MG/5ML;167MG/5ML	A080013 002	
--------------	-------------------------------	-------------	--

TERFONYL

BRISTOL MYERS SQUIBB	167MG/5ML;167MG/5ML;167MG/5ML	N006904 002	
----------------------	-------------------------------	-------------	--

TRIPLE SULFA

ALPHARMA US PHARMS	167MG/5ML;167MG/5ML;167MG/5ML	A080280 001	
--------------------	-------------------------------	-------------	--

TRIPLE SULFAS

LEDERLE	167MG/5ML;167MG/5ML;167MG/5ML	N006920 003	
---------	-------------------------------	-------------	--

TABLET;ORAL

NEOTRIZINE

LILLY	167MG;167MG;167MG	N006317 011	
-------	-------------------	-------------	--

SULFA-TRIPLE #2

IMPAX LABS	167MG;167MG;167MG	A080079 001	
------------	-------------------	-------------	--

SULFALOID

FOREST PHARMS	167MG;167MG;167MG	A080099 001	
---------------	-------------------	-------------	--

SULFOSE

WYETH AYERST	167MG;167MG;167MG	A080013 001	
--------------	-------------------	-------------	--

TERFONYL

BRISTOL MYERS SQUIBB	167MG;167MG;167MG	N006904 001	
----------------------	-------------------	-------------	--

TRIPLE SULFA

PUREPAC PHARM	167MG;167MG;167MG	A080086 001	
---------------	-------------------	-------------	--

TRIPLE SULFAS

LEDERLE	167MG;167MG;167MG	N006920 002	
---------	-------------------	-------------	--

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

TRISULFAPYRIMIDINES (SULFADIAZINE;SULFAMERAZINE;SULFAMETHAZINE)

TABLET;ORAL

TRIPLE SULFOID

PAL PAK

167MG;167MG;167MG

A080094 001

TROGLITAZONE

TABLET;ORAL

PRELAY

SANKYO

200MG

N020719 001 Jan 29, 1997

300MG

N020719 003 Aug 04, 1997

400MG

N020719 002 Jan 29, 1997

REZULIN

PFIZER PHARMS

200MG

N020720 001 Jan 29, 1997

300MG

N020720 003 Aug 04, 1997

400MG

N020720 002 Jan 29, 1997

TROLAMINE POLYPEPTIDE OLEATE CONDENSATE

SOLUTION/DROPS;OTIC

CERUMENEX

PHARM RES ASSOC

10%

N011340 002

TROLEANDOMYCIN

CAPSULE;ORAL

TAO

PFIZER

EQ 250MG BASE

N050336 002

SUSPENSION;ORAL

TAO

PFIZER

EQ 125MG BASE/5ML

N050332 001

TROPICAMIDE

SOLUTION/DROPS;OPHTHALMIC

MYDRIACYL

ALCON

0.5% **

N012111 002

1% **

N012111 004

MYDRIAFAIR

PHARMAFAIR

0.5%

A088274 001 Sep 16, 1983

1%

A088230 001 Sep 16, 1983

TROPICAMIDE

ALCON PHARMS LTD

1%

A089172 001 Dec 28, 1990

EPIC PHARMA LLC

1%

A088447 001 Aug 28, 1985

MIZA PHARMS USA

0.5%

A087636 001 Jul 30, 1982

1%

A087637 001 Aug 09, 1982

WATSON LABS

0.5%

A089171 001 Dec 28, 1990

TROSPIUM CHLORIDE

CAPSULE, EXTENDED RELEASE;ORAL

SANCTURA XR

+ ALLERGAN

60MG **

N022103 001 Aug 03, 2007

TROSPIUM CHLORIDE

UPSHER SMITH LABS

60MG

A091635 001 Apr 29, 2015

TABLET;ORAL

SANCTURA

+ ALLERGAN

20MG **

N021595 001 May 28, 2004

TROVAFLOXACIN MESYLATE

TABLET;ORAL

TROVAN

PFIZER

EQ 100MG BASE

N020759 001 Dec 18, 1997

EQ 200MG BASE

N020759 002 Dec 18, 1997

TRYPAN BLUE

SOLUTION;OPHTHALMIC

MEMBRANEBLUE

+ DORC

0.15%

N022278 001 Feb 20, 2009

TUBOCURARINE CHLORIDE

INJECTABLE;INJECTION

TUBOCURARINE CHLORIDE

BRISTOL MYERS SQUIBB

3MG/ML

N005657 001

HOSPIRA

3MG/ML

N006095 001

LILLY

3MG/ML

N006325 001

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

TYROPANOATE SODIUM

CAPSULE;ORAL

BILOPAQUE

GE HEALTHCARE

750MG

N013731 001

UMBRALISIB TOSYLATE

TABLET;ORAL

UKONIQ

+ TG THERAPS

EQ 200MG BASE

N213176 001 Feb 05, 2021

UNOPROSTONE ISOPROPYL

SOLUTION/DROPS;OPHTHALMIC

RESCULA

+ SUCAMPO PHARMA LLC

0.15% **

N021214 001 Aug 03, 2000

URACIL MUSTARD

CAPSULE;ORAL

URACIL MUSTARD

SHIRE

1MG

N012892 001

UREA

INJECTABLE;INJECTION

STERILE UREA

HOSPIRA

40GM/VIAL

N017698 001

UREAPHIL

HOSPIRA

40GM/VIAL

N012154 001

UREA C-13

FOR SOLUTION;ORAL

BREATHTEK UBT FOR H-PYLORI

OTSUKA AMERICA

EQ 75MG/POUCH

N020586 002 May 10, 2001

HELICOSOL

METABOLIC SOLUTIONS

125MG/VIAL

N021092 001 Dec 17, 1999

MERETEK UBT KIT (W/ PRANACTIN)

OTSUKA AMERICA

125MG/VIAL

N020586 001 Sep 17, 1996

PYLORI-CHEK BREATH TEST

DXS DEVICES

100MG/VIAL

N020900 001 Feb 04, 1999

UREA, C-14

CAPSULE;ORAL

PYTEST

+ AVENT

1uCi

N020617 001 May 09, 1997

PYTEST KIT

+ AVENT

1uCi **

N020617 002 May 09, 1997

URSODIOL

CAPSULE;ORAL

ACTIGALL

TEVA BRANDED PHARM

150MG

N019594 001 Dec 31, 1987

URSODIOL

IMPAX LABS INC

300MG

A077895 001 Jul 27, 2006

NORVIUM BIOSCIENCE

300MG

A090530 001 Feb 17, 2010

TEVA PHARMS

300MG

A075592 001 May 25, 2000

TABLET;ORAL

URSODIOL

IMPAX LABS INC

250MG

A200826 001 Dec 23, 2011

500MG

A200826 002 Dec 23, 2011

TEVA PHARMS USA

250MG

A079184 001 May 13, 2009

500MG

A079184 002 May 13, 2009

VALACYCLOVIR HYDROCHLORIDE

TABLET;ORAL

VALACYCLOVIR HYDROCHLORIDE

CIPLA

EQ 500MG BASE

A077135 001 May 24, 2010

EQ 1GM BASE

A077135 002 May 24, 2010

HIKMA

EQ 500MG BASE

A078656 001 May 24, 2010

EQ 1GM BASE

A078656 002 May 24, 2010

NORVIUM BIOSCIENCE

EQ 500MG BASE

A078070 001 May 24, 2010

EQ 1GM BASE

A078070 002 May 24, 2010

SANDOZ

EQ 500MG BASE

A077478 001 May 24, 2010

EQ 1GM BASE

A077478 002 May 24, 2010

TEVA PHARMS

EQ 500MG BASE

A077655 001 May 24, 2010

EQ 1GM BASE

A077655 002 May 24, 2010

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

VALACYCLOVIR HYDROCHLORIDE

TABLET; ORAL

VALACYCLOVIR HYDROCHLORIDE

WATSON LABS INC

EQ 500MG BASE

A090370 001 Mar 16, 2011

EQ 1GM BASE

A090370 002 Mar 16, 2011

VALDECOXIB

TABLET; ORAL

BEXTRA

GD SEARLE

10MG

N021341 002 Nov 16, 2001

20MG

N021341 003 Nov 16, 2001

VALGANCICLOVIR HYDROCHLORIDE

TABLET; ORAL

VALGANCICLOVIR HYDROCHLORIDE

CIPLA

EQ 450MG BASE

A209672 001 Nov 09, 2018

NORVIUM BIOSCIENCE

EQ 450MG BASE

A205151 001 Mar 03, 2021

VALPROATE SODIUM

INJECTABLE; INJECTION

DEPACON

+ ABBVIE

EQ 100MG BASE/ML **

N020593 001 Dec 30, 1996

VALPROATE SODIUM

NORVIUM BIOSCIENCE

EQ 100MG BASE/ML

A208120 001 Dec 22, 2021

VALPROIC ACID

CAPSULE; ORAL

DEPAKENE

+ ABBVIE

250MG **

N018081 001

VALPROIC ACID

HIBROW HLTHCARE

250MG

A207611 001 Aug 05, 2019

PAR PHARM

250MG

A070431 001 Feb 28, 1986

SCHERER RP

250MG

A070195 001 Jul 02, 1987

SUN PHARM INDS LTD

250MG

A091037 001 Feb 22, 2013

UPSHER SMITH LABS

250MG

A070631 001 Jun 11, 1987

CAPSULE, DELAYED RELEASE; ORAL

STAVZOR

+ BIONPHARMA

125MG **

N022152 001 Jul 29, 2008

+

250MG **

N022152 002 Jul 29, 2008

+

500MG **

N022152 003 Jul 29, 2008

SYRUP; ORAL

DEPAKENE

+ ABBVIE

250MG/5ML **

N018082 001

VALPROIC ACID

HIKMA

250MG/5ML

A074060 001 Jan 13, 1995

LANNETT CO INC

250MG/5ML

A077960 001 Oct 13, 2006

NOSTRUM LABS INC

250MG/5ML

A077105 001 Jul 29, 2005

PHARMOBEDIANT CNSLTG

250MG/5ML

A070868 001 Jul 01, 1986

VALSARTAN

CAPSULE; ORAL

DIOVAN

NOVARTIS

80MG

N020665 001 Dec 23, 1996

160MG

N020665 002 Dec 23, 1996

SOLUTION; ORAL

PREXXARTAN

+ CARMEL BIOSCIENCES

20MG/5ML **

N209139 001 Dec 19, 2017

+

80MG/20ML **

N209139 002 Dec 19, 2017

TABLET; ORAL

VALSARTAN

IVAX PHARMS

40MG

A077530 001 Jan 04, 2016

80MG

A077530 002 Jan 04, 2016

160MG

A077530 003 Jan 04, 2016

320MG

A077530 004 Jan 04, 2016

TORRENT

40MG

A202728 001 Jan 05, 2015

80MG

A202728 002 Jan 05, 2015

160MG

A202728 003 Jan 05, 2015

320MG

A202728 004 Jan 05, 2015

UNICHEM

40MG

A209261 001 May 04, 2018

80MG

A209261 002 May 04, 2018

160MG

A209261 003 May 04, 2018

320MG

A209261 004 May 04, 2018

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

VALSARTAN

TABLET; ORAL

VALSARTAN

WATSON LABS INC	40MG	A090642 001	Jan 05, 2015
	80MG	A090642 002	Jan 05, 2015
	160MG	A090642 003	Jan 05, 2015
	320MG	A090642 004	Jan 05, 2015

VANCOMYCIN

SOLUTION; INTRAVENOUS, ORAL

VANCOMYCIN

+ HIKMA	5GM/100ML (50MG/ML)	N213895 001	Aug 26, 2021
---------	---------------------	-------------	--------------

VANCOMYCIN HYDROCHLORIDE

CAPSULE; ORAL

VANCOMYCIN HYDROCHLORIDE

FRESENIUS KABI USA	EQ 125MG BASE	A065453 001	Jun 18, 2012
	EQ 250MG BASE	A065453 002	Jun 18, 2012
PAI HOLDINGS PHARM	EQ 125MG BASE	A065478 001	Apr 09, 2012
	EQ 250MG BASE	A065478 002	Apr 09, 2012
WATSON LABS	EQ 125MG BASE	A065510 001	Apr 09, 2012
	EQ 250MG BASE	A065510 002	Apr 09, 2012

FOR SOLUTION; ORAL

VANCOCIN HYDROCHLORIDE

ANI PHARMS	EQ 500MG BASE/6ML	A061667 001	
------------	-------------------	-------------	--

VANCOLED

LEDERLE	EQ 250MG BASE/5ML	A063321 002	Oct 15, 1993
	EQ 500MG BASE/6ML	A063321 003	Oct 15, 1993

INJECTABLE; INJECTION

VANCOCIN HYDROCHLORIDE

STERISCIENCE SPECLTS	EQ 500MG BASE/VIAL **	A060180 001	
	EQ 500MG BASE/VIAL	A062476 001	Mar 15, 1984
	EQ 500MG BASE/VIAL	A062716 001	Mar 13, 1987
	EQ 500MG BASE/VIAL **	A062812 001	Nov 17, 1987
	EQ 1GM BASE/VIAL **	A060180 002	Mar 21, 1986
	EQ 1GM BASE/VIAL	A062476 002	Mar 21, 1986
	EQ 1GM BASE/VIAL	A062716 002	Mar 13, 1987
	EQ 1GM BASE/VIAL **	A062812 002	Nov 17, 1987
	EQ 10GM BASE/VIAL **	A062812 003	Nov 17, 1987

VANCOLED

HIKMA	EQ 500MG BASE/VIAL **	A062682 001	Jul 22, 1986
	EQ 1GM BASE/VIAL **	A062682 002	Mar 30, 1988
	EQ 2GM BASE/VIAL **	A062682 003	May 11, 1988
	EQ 5GM BASE/VIAL **	A062682 004	May 11, 1988
	EQ 10GM BASE/VIAL **	A062682 005	May 11, 1988

VANCOMYCIN HYDROCHLORIDE

AVET LIFESCIENCES	EQ 500MG BASE/VIAL	A202275 001	Oct 31, 2013
	EQ 1GM BASE/VIAL	A202275 002	Oct 31, 2013
	EQ 10GM BASE/VIAL	A202464 001	Oct 09, 2013
	EQ 5GM BASE/VIAL	A202274 001	Oct 31, 2013
HIKMA	EQ 500MG BASE/VIAL	A062879 001	Aug 02, 1988
	EQ 500MG BASE/VIAL	A203300 001	Aug 11, 2020
	EQ 1GM BASE/VIAL	A062879 002	Aug 02, 1988
KNACK	EQ 500MG BASE/VIAL	A213059 001	Feb 15, 2022
	EQ 750MG BASE/VIAL	A213059 002	Feb 15, 2022
	EQ 1GM BASE/VIAL	A213059 003	Feb 15, 2022
MEDIMETRIKS PHARMS	EQ 1GM BASE/VIAL	A065401 002	Jun 30, 2008
	EQ 500MG BASE/VIAL	A065401 001	Jun 30, 2008
MYLAN LABS LTD	EQ 10GM BASE/VIAL	A091469 001	Jul 01, 2011
SANDOZ	EQ 500MG BASE/VIAL	A090250 001	Apr 27, 2010
	EQ 1GM BASE/VIAL	A090250 002	Apr 27, 2010
SANDOZ INC	EQ 5GM BASE/VIAL	A201048 001	Aug 10, 2012
	EQ 10GM BASE/VIAL	A201048 002	Aug 10, 2012
TEVA PHARMS USA	EQ 500MG BASE/VIAL	A201251 001	Dec 23, 2015
	EQ 1GM BASE/VIAL	A201251 002	Dec 23, 2015
	EQ 5GM BASE/VIAL	A201250 001	Dec 23, 2015
	EQ 10GM BASE/VIAL	A201250 002	Dec 23, 2015
XELLIA PHARMS APS	EQ 500MG BASE/VIAL	A091377 001	Sep 09, 2015
	EQ 1GM BASE/VIAL	A091377 002	Sep 09, 2015
	EQ 5GM BASE/VIAL	A206243 001	Dec 23, 2015

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

VANCOMYCIN HYDROCHLORIDE

INJECTABLE; INJECTION

VANCOMYCIN HYDROCHLORIDE

EQ 10GM BASE/VIAL

A206243 002 Dec 23, 2015

VANCOR

PHARMACIA AND UPJOHN

EQ 500MG BASE/VIAL

A062956 001 Aug 01, 1988

EQ 1GM BASE/VIAL

A062956 002 Aug 01, 1988

POWDER; INTRAVENOUS

VANCOMYCIN HYDROCHLORIDE

+ MYLAN LABS LTD

EQ 250MG BASE/VIAL **

N209481 001 Jul 10, 2018

VARDENAFIL HYDROCHLORIDE

TABLET; ORAL

LEVITRA

+ BAYER HLTHCARE

EQ 2.5MG BASE **

N021400 003 Aug 19, 2003

+

EQ 5MG BASE **

N021400 001 Aug 19, 2003

+

EQ 10MG BASE **

N021400 002 Aug 19, 2003

+

EQ 20MG BASE **

N021400 004 Aug 19, 2003

VARDENAFIL HYDROCHLORIDE

STEVENS J

EQ 5MG BASE

A210738 001 Oct 31, 2018

EQ 10MG BASE

A210738 002 Oct 31, 2018

EQ 20MG BASE

A210738 003 Oct 31, 2018

TABLET, ORALLY DISINTEGRATING; ORAL

STAXYN

+ BAYER HLTHCARE

10MG **

N200179 001 Jun 17, 2010

VARENICLINE TARTRATE

TABLET; ORAL

CHANTIX

+ PF PRISM CV

EQ 0.5MG BASE **

N021928 001 May 10, 2006

+

EQ 1MG BASE **

N021928 002 May 10, 2006

VASOPRESSIN

SOLUTION; INTRAVENOUS

VASOSTRICT

+ ENDO OPERATIONS

50UNITS/50ML (1UNITS/ML)

N204485 006 Apr 12, 2023

+

60UNITS/100ML (0.6UNITS/ML)

N204485 004 Apr 15, 2020

VASOPRESSIN TANNATE

INJECTABLE; INJECTION

PITRESSIN TANNATE

+ PARKE DAVIS

5PRESSOR UNITS/ML **

N003402 001

VECURONIUM BROMIDE

INJECTABLE; INJECTION

NORCURON

+ ORGANON USA INC

10MG/VIAL **

N018776 002 Apr 30, 1984

+

20MG/VIAL **

N018776 003 Jan 03, 1992

VECURONIUM BROMIDE

HIKMA

10MG/VIAL

A075218 001 Aug 23, 1999

20MG/VIAL

A075218 002 Aug 23, 1999

HOSPIRA

4MG/VIAL

A075558 001 Sep 11, 2001

WATSON LABS

10MG/VIAL

A074334 001 Aug 31, 1995

20MG/VIAL

A074334 002 Aug 31, 1995

VENLAFAXINE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

EFFEXOR XR

UPJOHN

EQ 100MG BASE

N020699 003 Oct 20, 1997

VENLAFAXINE HYDROCHLORIDE

ANCHEN PHARMS

EQ 37.5MG BASE

A078087 001 Mar 16, 2012

EQ 75MG BASE

A078087 002 Mar 16, 2012

EQ 150MG BASE

A078087 003 Mar 16, 2012

NORVIUM BIOSCIENCE

EQ 37.5MG BASE

A078789 001 Jun 01, 2011

EQ 75MG BASE

A078789 002 Jun 01, 2011

EQ 150MG BASE

A078789 003 Jun 01, 2011

NOSTRUM PHARMS LLC

EQ 37.5MG BASE

A200430 001 Apr 04, 2023

EQ 75MG BASE

A200430 002 Apr 04, 2023

EQ 150MG BASE

A200430 003 Apr 04, 2023

TORRENT

EQ 37.5MG BASE

A090899 001 Jun 01, 2011

EQ 75MG BASE

A090899 002 Jun 01, 2011

EQ 150MG BASE

A090899 003 Jun 01, 2011

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

VENLAFAXINE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE;ORAL

VENLAFAXINE HYDROCHLORIDE

VALEANT PHARMS NORTH	EQ 37.5MG BASE	A090071 001	Apr 15, 2011
	EQ 75MG BASE	A090071 002	Apr 15, 2011
	EQ 150MG BASE	A090071 003	Apr 15, 2011

TABLET;ORAL

EFFEXOR

+	WYETH PHARMS INC	EQ 12.5MG BASE **	N020151 001	Dec 28, 1993
+		EQ 25MG BASE **	N020151 002	Dec 28, 1993
+		EQ 37.5MG BASE **	N020151 006	Dec 28, 1993
+		EQ 50MG BASE **	N020151 003	Dec 28, 1993
+		EQ 75MG BASE **	N020151 004	Dec 28, 1993
+		EQ 100MG BASE **	N020151 005	Dec 28, 1993

VENLAFAXINE HYDROCHLORIDE

AMNEAL PHARMS

EQ 25MG BASE	A079098 001	May 11, 2010
EQ 37.5MG BASE	A079098 002	May 11, 2010
EQ 50MG BASE	A079098 003	May 11, 2010
EQ 75MG BASE	A079098 004	May 11, 2010
EQ 100MG BASE	A079098 005	May 11, 2010

CHARTWELL RX

EQ 25MG BASE	A077515 001	Jun 13, 2008
EQ 37.5MG BASE	A077515 002	Jun 13, 2008
EQ 50MG BASE	A077515 003	Jun 13, 2008
EQ 75MG BASE	A077515 004	Jun 13, 2008
EQ 100MG BASE	A077515 005	Jun 13, 2008

NORVIUM BIOSCIENCE

EQ 25MG BASE	A077166 001	Jun 13, 2008
EQ 37.5MG BASE	A077166 002	Jun 13, 2008
EQ 50MG BASE	A077166 003	Jun 13, 2008
EQ 75MG BASE	A077166 004	Jun 13, 2008
EQ 100MG BASE	A077166 005	Jun 13, 2008

PLIVA HRVATSKA DOO

EQ 25MG BASE	A078517 001	Jun 13, 2008
EQ 37.5MG BASE	A078517 002	Jun 13, 2008
EQ 50MG BASE	A078517 003	Jun 13, 2008
EQ 75MG BASE	A078517 004	Jun 13, 2008
EQ 100MG BASE	A078517 005	Jun 13, 2008

PRINSTON INC

EQ 25MG BASE	A090027 001	Aug 04, 2010
EQ 37.5MG BASE	A090027 002	Aug 04, 2010
EQ 50MG BASE	A090027 003	Aug 04, 2010
EQ 75MG BASE	A090027 004	Aug 04, 2010
EQ 100MG BASE	A090027 005	Aug 04, 2010

TABLET, EXTENDED RELEASE;ORAL

VENLAFAXINE HYDROCHLORIDE

SUN PHARM

EQ 37.5MG BASE	A091272 001	Aug 18, 2010
EQ 75MG BASE	A091272 002	Aug 18, 2010
EQ 150MG BASE	A091272 003	Aug 18, 2010
EQ 225MG BASE	A091272 004	Jan 08, 2019

SWISS PHARM

EQ 75MG BASE	A214423 001	Jan 04, 2022
EQ 150MG BASE	A214423 002	Jan 04, 2022

VERAPAMIL HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE;ORAL

VERAPAMIL HYDROCHLORIDE

RISING

100MG	A078306 001	Aug 09, 2007
200MG	A078306 002	Aug 09, 2007
300MG	A078306 003	Aug 09, 2007

INJECTABLE;INJECTION

CALAN

GD SEARLE LLC

2.5MG/ML	N019038 001	Mar 30, 1984
----------	-------------	--------------

ISOPTIN

+ MT ADAMS

2.5MG/ML **	N018485 001	
-------------	-------------	--

VERAPAMIL HYDROCHLORIDE

ABRAXIS PHARM

2.5MG/ML	A070348 001	May 01, 1986
----------	-------------	--------------

BEDFORD

2.5MG/ML	A072888 001	Jul 28, 1995
----------	-------------	--------------

HOSPIRA

2.5MG/ML	A070577 001	Feb 02, 1987
----------	-------------	--------------

2.5MG/ML	A070739 001	May 06, 1987
----------	-------------	--------------

2.5MG/ML	A070740 001	May 06, 1987
----------	-------------	--------------

INTL MEDICATION

2.5MG/ML	A070451 001	Dec 16, 1985
----------	-------------	--------------

LUITPOLD

2.5MG/ML	A070225 001	Nov 12, 1985
----------	-------------	--------------

2.5MG/ML	A070617 001	Nov 12, 1985
----------	-------------	--------------

MARSAM PHARMS LLC

2.5MG/ML	A072233 001	Feb 26, 1993
----------	-------------	--------------

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

VERAPAMIL HYDROCHLORIDE

INJECTABLE; INJECTION

VERAPAMIL HYDROCHLORIDE

	2.5MG/ML		A073485	001	Sep 27, 1993
SMITH AND NEPHEW	2.5MG/ML		A070696	001	Jul 31, 1987
	2.5MG/ML		A070697	001	Jul 31, 1987
SOLOPAK	2.5MG/ML		A070695	001	Jul 31, 1987

SOLUTION; INTRAVENOUS

VERAPAMIL HYDROCHLORIDE

EXELA PHARMA	10MG/4ML (2.5MG/ML)		N018925	002	Apr 05, 2018
--------------	---------------------	--	---------	-----	--------------

TABLET; ORAL

CALAN

PFIZER	40MG **		N018817	003	Feb 23, 1988
+	80MG **		N018817	001	Sep 10, 1984
+	120MG **		N018817	002	Sep 10, 1984
	160MG **		N018817	004	Feb 23, 1988

ISOPTIN

MT ADAMS	40MG **		N018593	003	Nov 23, 1987
+	80MG **		N018593	001	Mar 08, 1982
+	120MG **		N018593	002	Mar 08, 1982

VERAPAMIL HYDROCHLORIDE

ACTAVIS ELIZABETH

	80MG		A071019	001	Sep 24, 1986
--	------	--	---------	-----	--------------

	120MG		A070468	001	Sep 24, 1986
--	-------	--	---------	-----	--------------

CHARTWELL RX	40MG		A073168	001	Jul 31, 1992
--------------	------	--	---------	-----	--------------

	80MG		A071423	001	May 24, 1988
--	------	--	---------	-----	--------------

	120MG		A071424	001	May 25, 1988
--	-------	--	---------	-----	--------------

MUTUAL PHARM	80MG		A070482	001	Sep 24, 1986
--------------	------	--	---------	-----	--------------

	120MG		A070483	001	Sep 24, 1986
--	-------	--	---------	-----	--------------

PLIVA	40MG		A072751	001	Feb 23, 1996
-------	------	--	---------	-----	--------------

	80MG		A072124	001	Jan 26, 1989
--	------	--	---------	-----	--------------

	120MG		A072125	001	Jan 26, 1989
--	-------	--	---------	-----	--------------

RISING	80MG		A071483	002	Feb 15, 1989
--------	------	--	---------	-----	--------------

	120MG		A071483	001	Feb 15, 1989
--	-------	--	---------	-----	--------------

SUN PHARM INDUSTRIES	80MG		A071489	002	Jan 13, 1988
----------------------	------	--	---------	-----	--------------

	120MG		A071489	001	Jan 13, 1988
--	-------	--	---------	-----	--------------

WARNER CHILCOTT	80MG		A070340	001	Sep 24, 1986
-----------------	------	--	---------	-----	--------------

	120MG		A070341	001	Sep 24, 1986
--	-------	--	---------	-----	--------------

WATSON LABS	40MG		A072799	001	Apr 28, 1989
-------------	------	--	---------	-----	--------------

	40MG		A072923	001	Jun 29, 1993
--	------	--	---------	-----	--------------

	80MG		A070855	001	Sep 24, 1986
--	------	--	---------	-----	--------------

	80MG		A071366	001	Oct 01, 1986
--	------	--	---------	-----	--------------

	120MG		A070856	001	Sep 24, 1986
--	-------	--	---------	-----	--------------

	120MG		A071367	001	Oct 01, 1986
--	-------	--	---------	-----	--------------

TABLET, EXTENDED RELEASE; ORAL

CALAN SR

PFIZER	120MG **		N019152	003	Mar 06, 1991
+	180MG **		N019152	002	Dec 15, 1989
+	240MG **		N019152	001	Dec 16, 1986

COVERA-HS

PFIZER	180MG		N020552	001	Feb 26, 1996
--------	-------	--	---------	-----	--------------

	240MG		N020552	002	Feb 26, 1996
--	-------	--	---------	-----	--------------

VERAPAMIL HYDROCHLORIDE

APOTEX CORP	120MG		A200878	001	Apr 20, 2012
-------------	-------	--	---------	-----	--------------

	180MG		A200878	002	Apr 20, 2012
--	-------	--	---------	-----	--------------

	240MG		A200878	003	Apr 20, 2012
--	-------	--	---------	-----	--------------

IVAX SUB TEVA PHARMS	120MG		A073568	002	Oct 10, 1997
----------------------	-------	--	---------	-----	--------------

	180MG		A074330	001	Jan 31, 1994
--	-------	--	---------	-----	--------------

	240MG		A073568	001	Jul 31, 1992
--	-------	--	---------	-----	--------------

PLIVA	240MG		A072922	001	Mar 01, 1996
-------	-------	--	---------	-----	--------------

RISING	120MG		A074587	002	Feb 21, 1997
--------	-------	--	---------	-----	--------------

	180MG		A074587	003	Sep 09, 1997
--	-------	--	---------	-----	--------------

	240MG		A074587	001	Mar 23, 1996
--	-------	--	---------	-----	--------------

SUN PHARM INDS INC	120MG		A090529	001	Dec 30, 2011
--------------------	-------	--	---------	-----	--------------

	180MG		A090529	002	Dec 30, 2011
--	-------	--	---------	-----	--------------

	240MG		A090529	003	Dec 30, 2011
--	-------	--	---------	-----	--------------

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

VERATRUM VIRIDE ROOT

TABLET; ORAL

VERTAVIS

MEDPOINTE PHARM HLC 130CSR UNIT N005691 002

VIDARABINE

INJECTABLE; INJECTION

VIRA-A

PARKEDALE EQ 187.4MG BASE/ML N050523 001

OINTMENT; OPHTHALMIC

VIRA-A

PARKEDALE 3% N050486 001

VIGABATRIN

FOR SOLUTION; ORAL

VIGABATRIN

ACCORD HLTHCARE 500MG/PACKET A214425 001 Nov 13, 2020

CHARTWELL RX 500MG/PACKET A211790 001 Mar 10, 2022

GRANULES 500MG/PACKET A213469 001 Apr 24, 2020

PROPEL PHARMA 500MG/PACKET A213390 001 Jul 29, 2021

SPECGX LLC 500MG/PACKET A212626 001 Jul 28, 2021

ZYDUS LIFESCIENCES 500MG/PACKET A214671 001 Mar 02, 2023

VINBLASTINE SULFATE

INJECTABLE; INJECTION

VELBAN

+ LILLY 10MG/VIAL ** N012665 001

VINBLASTINE SULFATE

ABRAXIS PHARM 10MG/VIAL A089011 001 Nov 18, 1985

HOSPIRA 10MG/VIAL A089565 001 Aug 18, 1987

VINCRIStINE SULFATE

INJECTABLE; INJECTION

ONCOVIN

+ LILLY 1MG/VIAL ** N014103 001

+ 1MG/ML ** N014103 003 Mar 07, 1984

+ 5MG/VIAL ** N014103 002

VINCASAR PFS

TEVA PARENTERAL 1MG/ML A071426 001 Jul 17, 1987

VINCREX

BRISTOL MYERS SQUIBB 5MG/VIAL A070867 001 Jul 12, 1988

VINCRIStINE SULFATE

ABIC 1MG/ML A070873 001 Feb 19, 1987

ABRAXIS PHARM 1MG/ML A070411 001 Sep 10, 1986

FRESENIUS KABI USA 1MG/ML A076296 001 Dec 20, 2002

1MG/ML A076401 001 Oct 28, 2003

HOSPIRA 1MG/VIAL A071559 001 Apr 11, 1988

2MG/VIAL A071560 001 Apr 11, 1988

5MG/VIAL A071561 001 Apr 11, 1988

VINCRIStINE SULFATE PFS

TEVA PHARMS USA 1MG/ML A075493 001 Sep 01, 1999

INJECTABLE, LIPOSOMAL; INTRAVENOUS

MARQIBO KIT

+ ACROTECH 5MG/5ML (1MG/ML) N202497 001 Aug 09, 2012

VINORELBINE TARTRATE

INJECTABLE; INJECTION

NAVELBINE

+ PIERRE FABRE EQ 10MG BASE/ML ** N020388 001 Dec 23, 1994

VINORELBINE TARTRATE

EBEWE PHARMA EQ 10MG BASE/ML A078408 001 Feb 13, 2008

FRESENIUS KABI USA EQ 10MG BASE/ML A076849 001 Apr 18, 2005

HOSPIRA EQ 10MG BASE/ML A076827 001 Jun 02, 2005

NOVAST LABS EQ 10MG BASE/ML A208997 001 Aug 05, 2019

RISING EQ 10MG BASE/ML A200148 001 Aug 31, 2012

TEVA PHARMS USA EQ 10MG BASE/ML A076028 001 Feb 03, 2003

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

VIOMYCIN SULFATE

INJECTABLE; INJECTION

VIOCIN SULFATE

PFIZER

EQ 1GM BASE/VIAL

A061086 001

EQ 5GM BASE/VIAL

A061086 002

VITAMIN A

CAPSULE; ORAL

AQUASOL A

ASTRAZENECA

25,000USP UNITS

A083080 002

50,000USP UNITS

A083080 001

VITAMIN A

BANNER PHARMACAPS

50,000USP UNITS

A083973 001

CHASE CHEM

50,000 IU

A083351 001

EVERYLIFE

50,000 IU

A083134 001

IMPAX LABS

50,000USP UNITS

A080952 001

WEST WARD

50,000USP UNITS

A080985 001

VITAMIN A PALMITATE

CAPSULE; ORAL

AFAXIN

STERLING WINTHROP

EQ 50,000 UNITS BASE

A083187 001

ALPHALIN

LILLY

EQ 50,000 UNITS BASE

A080883 001

DEL-VI-A

DEL RAY LABS

EQ 50,000 UNITS BASE

A080830 001

VI-DOM-A

BAYER PHARMS

EQ 50,000 UNITS BASE

A080972 001

VITAMIN A

BANNER PHARMACAPS

EQ 50,000 UNITS BASE

A080702 001

BRISTOL MYERS SQUIBB

EQ 50,000 UNITS BASE

A080860 001

CHASE CHEM

EQ 50,000 UNITS BASE

A080746 001

EQ 50,000 UNITS BASE

A083207 001

ELKINS SINN

EQ 50,000 UNITS BASE

A085479 001

EVERYLIFE

EQ 50,000 UNITS BASE

A080943 001

EQ 50,000 UNITS BASE

A083114 001

IMPAX LABS

EQ 50,000 UNITS BASE

A080953 001

EQ 50,000 UNITS BASE

A080955 001

IVAX SUB TEVA PHARMS

EQ 50,000 UNITS BASE

A083035 001

EQ 50,000 UNITS BASE

A083190 001

MK LABS

EQ 25,000 UNITS BASE

A083457 002

EQ 50,000 UNITS BASE

A083457 001

WEST WARD

EQ 50,000 UNITS BASE

A080967 001

WHARTON LABS

EQ 50,000 UNITS BASE

A083665 001

VITAMIN A PALMITATE

ARCUM

EQ 50,000 UNITS BASE

A083311 001

EQ 50,000 UNITS BASE

A083321 001

BANNER PHARMACAPS

EQ 50,000 UNITS BASE

A083948 001

EQ 50,000 UNITS BASE

A083981 001

VITAMIN A SOLUBILIZED

TEVA

EQ 50,000 UNITS BASE

A080921 001

INJECTABLE; INJECTION

VITAMIN A PALMITATE

BEL MAR

EQ 50,000 UNITS BASE/ML

A080819 001

VORAPAXAR SULFATE

TABLET; ORAL

ZONTIVITY

+ KEY THERAP

EQ 2.08MG BASE

N204886 001 May 08, 2014

VORICONAZOLE

FOR SUSPENSION; ORAL

VORICONAZOLE

RISING

200MG/5ML

A202361 001 May 28, 2013

INJECTABLE; INTRAVENOUS

VORICONAZOLE

EUGIA PHARMA

200MG/VIAL

A212162 001 Feb 02, 2023

NORVIUM BIOSCIENCE

200MG/VIAL

A210849 001 Oct 11, 2022

TABLET; ORAL

VORICONAZOLE

TEVA PHARMS

50MG

A091658 001 Apr 06, 2012

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

VORICONAZOLETABLET; ORAL
VORICONAZOLE

200MG

A091658 002 Apr 06, 2012

VORTIOXETINE HYDROBROMIDETABLET; ORAL
TRINTELLIX

+ TAKEDA PHARMS USA EQ 15MG BASE **

N204447 003 Sep 30, 2013

VOXELOTOR

TABLET; ORAL

OXBRYTA

+ GLOBAL BLOOD THERAPS 300MG

N213137 002 Oct 14, 2022

+ 500MG

N213137 001 Nov 25, 2019

TABLET, FOR SUSPENSION; ORAL

OXBRYTA

+ GLOBAL BLOOD THERAPS 300MG

N216157 001 Dec 17, 2021

WARFARIN POTASSIUM

TABLET; ORAL

ATHROMBIN-K

PHARM RES ASSOC

2MG

N011771 007

5MG

N011771 004

10MG

N011771 005

25MG

N011771 006

WARFARIN SODIUM

INJECTABLE; INJECTION

COUMADIN

BRISTOL MYERS SQUIBB 5MG/VIAL

N009218 024 Feb 07, 1995

50MG/VIAL

N009218 020

75MG/VIAL

N009218 012

TABLET; ORAL

ATHROMBIN

PHARM RES ASSOC

5MG

N011771 003

10MG

N011771 002

25MG

N011771 001

COUMADIN

+ BRISTOL MYERS SQUIBB 1MG **

N009218 022 Mar 01, 1990

+ 2MG

N009218 013

+ 2.5MG

N009218 018

+ 3MG

N009218 025 Nov 18, 1996

+ 4MG

N009218 023 Aug 24, 1993

+ 5MG

N009218 007

+ 6MG

N009218 026 Nov 18, 1996

+ 7.5MG

N009218 016

+ 10MG

N009218 005

PANWARFIN

ABBOTT

2MG

N017020 001

2.5MG

N017020 002

5MG

N017020 003

7.5MG

N017020 004

10MG

N017020 005

WARFARIN SODIUM

AUROBINDO PHARMA USA

1MG

A040415 001 Sep 27, 2004

2MG

A040415 002 Sep 27, 2004

2.5MG

A040415 003 Sep 29, 2004

3MG

A040415 004 Sep 27, 2004

4MG

A040415 005 Sep 27, 2004

5MG

A040415 006 Sep 27, 2004

6MG

A040415 007 Sep 27, 2004

7.5MG

A040415 008 Sep 27, 2004

10MG

A040415 009 Sep 27, 2004

BARR

1MG

A040145 001 Mar 26, 1997

2MG

A040145 002 Mar 26, 1997

2.5MG

A040145 003 Mar 26, 1997

3MG

A040145 008 Nov 05, 1998

4MG

A040145 004 Mar 26, 1997

5MG

A040145 005 Mar 26, 1997

6MG

A040145 009 Nov 05, 1998

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

WARFARIN SODIUM

TABLET; ORAL

WARFARIN SODIUM

	7.5MG	A040145 006	Mar 26, 1997
	10MG	A040145 007	Mar 26, 1997
CHARTWELL RX	1MG	A040196 001	Sep 30, 1997
	2MG	A040196 002	Sep 30, 1997
	2.5MG	A040196 003	Sep 30, 1997
	3MG	A040196 008	Jul 26, 2000
	4MG	A040196 004	Sep 30, 1997
	5MG	A040196 005	Sep 30, 1997
	6MG	A040196 009	Jul 26, 2000
	7.5MG	A040196 006	Sep 30, 1997
	10MG	A040196 007	Sep 30, 1997
IPCA LABS LTD	1MG	A200104 001	Jun 27, 2013
	2MG	A200104 002	Jun 27, 2013
	2.5MG	A200104 003	Jun 27, 2013
	3MG	A200104 004	Jun 27, 2013
	4MG	A200104 005	Jun 27, 2013
	5MG	A200104 006	Jun 27, 2013
	6MG	A200104 007	Jun 27, 2013
	7.5MG	A200104 008	Jun 27, 2013
	10MG	A200104 009	Jun 27, 2013
USL PHARMA	2MG	A088719 001	Jun 27, 1985
	2.5MG	A088720 001	Aug 06, 1985
	5MG	A088721 001	Jul 02, 1985
WATSON LABS	2MG	A086123 001	Aug 17, 1982
	2.5MG	A086120 001	Aug 17, 1982
	5MG	A086119 001	Aug 17, 1982
	7.5MG	A086118 001	Aug 17, 1982
	10MG	A086122 001	Aug 17, 1982

XENON XE-127

GAS; INHALATION

XENON XE 127

MALLINCKRODT	5mCi/VIAL	N018536 001	Oct 01, 1982
	10mCi/VIAL	N018536 002	Oct 01, 1982

XENON XE-133

GAS; INHALATION

XENON XE 133

GE HEALTHCARE	1 CI/AMP	N017256 002	
	10mCi/VIAL	N017687 002	
	20mCi/VIAL	N017687 003	
GEN ELECTRIC	5-100 CI/CYLINDER	N017550 001	
	0.25-5 CI/AMP	N017550 003	
XENON XE 133-V.S.S.			
GE HEALTHCARE	10mCi/VIAL	N017687 001	
INJECTABLE; INJECTION			
XENON XE 133			
GE HEALTHCARE	1.3-1.7 CI/AMP	N017256 001	
LANTHEUS MEDCL	6.3mCi/ML	N017283 001	
SOLUTION; INHALATION, INJECTION			
XENEISOL			
MALLINCKRODT	18-25mCi/AMP	N017262 002	

XYLOSE

POWDER; ORAL

XYLO-PFAN

SAVAGE LABS	25GM/BOT	N017605 001	
XYLOSE			
LYNE	25GM/BOT	N018856 001	Mar 26, 1987

ZALCITABINE

TABLET; ORAL

HIVID

ROCHE	0.375MG	N020199 001	Jun 19, 1992
	0.75MG	N020199 002	Jun 19, 1992

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ZALEPLON

CAPSULE; ORAL

ZALEPLON

HIKMA PHARMS	5MG	A078147 001	Nov 25, 2008
	10MG	A078147 002	Nov 25, 2008
NATCO PHARMA	5MG	A077238 001	Jun 06, 2008
	10MG	A077238 002	Jun 06, 2008
TEVA PHARMS	5MG	A077239 001	Jun 06, 2008
	10MG	A077239 002	Jun 06, 2008
UPSHER SMITH LABS	5MG	A078095 001	Jun 06, 2008
	5MG	A078706 001	Jun 06, 2008
	10MG	A078095 002	Jun 06, 2008
	10MG	A078706 002	Jun 06, 2008

ZICONOTIDE ACETATE

INJECTABLE; INTRATHECAL

PRIALT

TERSERA	200MCG/2ML (100MCG/ML)	N021060 003	Dec 28, 2004
---------	------------------------	-------------	--------------

ZIDOVUDINE

INJECTABLE; INJECTION

ZIDOVUDINE

AM REGENT	10MG/ML	A091457 001	May 06, 2010
LIAONING CHENGDA	10MG/ML	A204538 001	Nov 26, 2013

TABLET; ORAL

RETROVIR

VIIV HLTHCARE	200MG	N020518 001	Dec 19, 1995
+	300MG **	N020518 002	Oct 04, 1996

ZIDOVUDINE

AUROBINDO PHARMA	60MG	N022294 001	Jul 23, 2009
HEC PHARM	300MG	A202058 001	Oct 07, 2011
HIKMA	300MG	A076844 001	Sep 19, 2005
NORVIUM BIOSCIENCE	100MG	N200732 001	Feb 23, 2011
	300MG	A078922 001	Feb 14, 2008
RANBAXY LABS LTD	300MG	A077327 001	Sep 19, 2005

ZILEUTON

TABLET; ORAL

ZYFLO

CHIESI	300MG	N020471 001	Dec 09, 1996
--------	-------	-------------	--------------

TABLET, EXTENDED RELEASE; ORAL

ZILEUTON

LUPIN LTD	600MG	A211972 001	Nov 05, 2019
TEVA PHARMS USA	600MG	A211043 001	May 03, 2022

ZYFLO CR

+	CHIESI	600MG **	N022052 001	May 30, 2007
---	--------	----------	-------------	--------------

ZINC SULFATE

INJECTABLE; INJECTION

ZINC SULFATE

ABRAXIS PHARM	EQ 1MG ZINC/ML	N019229 002	May 05, 1987
---------------	----------------	-------------	--------------

ZIPRASIDONE HYDROCHLORIDE

CAPSULE; ORAL

ZIPRASIDONE HYDROCHLORIDE

MYLAN	EQ 20MG BASE	A202395 001	Oct 10, 2013
	EQ 40MG BASE	A202395 002	Oct 10, 2013
	EQ 60MG BASE	A202395 003	Oct 10, 2013
	EQ 80MG BASE	A202395 004	Oct 10, 2013

SUSPENSION; ORAL

GEODON

PFIZER INC	EQ 10MG BASE/ML	N021483 001	Mar 29, 2006
------------	-----------------	-------------	--------------

ZOLEDRONIC ACID

INJECTABLE; INTRAVENOUS

ZOLEDRONIC ACID

ACTAVIS INC	EQ 4MG BASE/5ML	A202472 001	Mar 04, 2013
AVET LIFESCIENCES	EQ 5MG BASE/100ML	A201801 001	Mar 29, 2013
BRECKENRIDGE	EQ 4MG BASE/5ML	A091170 001	Mar 04, 2013
DR REDDYS	EQ 4MG BASE/100ML	A204344 001	Nov 19, 2018
EUGIA PHARMA	EQ 4MG BASE/5ML	A207751 001	Sep 26, 2016
	EQ 5MG BASE/100ML	A209125 001	Dec 08, 2017

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ZOLEDRONIC ACID

INJECTABLE; INTRAVENOUS

ZOLEDRONIC ACID

HOSPIRA	EQ 4MG BASE/5ML	A090621	001	Mar 19, 2015
NOVAST LABS	EQ 4MG BASE/5ML	A208968	001	Feb 19, 2020
SHILPA	EQ 4MG BASE/5ML	A208513	001	May 15, 2019
SUN PHARMA GLOBAL	EQ 4MG BASE/VIAL	A090018	001	Mar 04, 2013
	EQ 4MG BASE/5ML	A202746	001	Mar 04, 2013

ZOMETA

+ NOVARTIS	EQ 4MG BASE/VIAL **	N021223	001	Aug 20, 2001
+	EQ 4MG BASE/5ML **	N021223	002	Mar 07, 2003
+	EQ 4MG BASE/100ML **	N021223	003	Jun 17, 2011

ZOLMITRIPTAN

TABLET; ORAL

ZOLMITRIPTAN

ANI PHARMS	2.5MG	A090861	001	Mar 04, 2014
	5MG	A090861	002	Mar 04, 2014
APOTEX INC	2.5MG	A202078	001	May 14, 2013
	5MG	A202078	002	May 14, 2013
INVAGEN PHARMS	2.5MG	A204284	001	Apr 09, 2014
	5MG	A204284	002	Apr 09, 2014
MACLEODS PHARMS LTD	2.5MG	A203772	001	Sep 30, 2015
	5MG	A203772	002	Sep 30, 2015
NATCO PHARMA USA	2.5MG	A203186	001	May 14, 2013
	5MG	A203186	002	May 14, 2013
SUN PHARMA GLOBAL	2.5MG	A203476	001	Nov 13, 2014
	5MG	A203476	002	Nov 13, 2014

ZOMIG

+ IPR	2.5MG	N020768	001	Nov 25, 1997
+	5MG	N020768	002	Nov 25, 1997

TABLET, ORALLY DISINTEGRATING; ORAL

ZOLMITRIPTAN

APOTEX INC	2.5MG	A202476	001	May 14, 2013
	5MG	A202476	002	May 14, 2013
MACLEODS PHARMS LTD	2.5MG	A204336	001	Oct 22, 2015
	5MG	A204336	002	Oct 22, 2015
RISING	2.5MG	A202855	001	Sep 20, 2019
	5MG	A202855	002	Sep 20, 2019

ZOMIG-ZMT

+ ASTRAZENECA	2.5MG	N021231	001	Feb 13, 2001
+	5MG	N021231	002	Sep 17, 2001

ZOLPIDEM TARTRATE

SPRAY, METERED; ORAL

ZOLPIMIST

+ AYTU	5MG/SPRAY	N022196	001	Dec 19, 2008
--------	-----------	---------	-----	--------------

TABLET; ORAL

ZOLPIDEM TARTRATE

DR REDDYS LABS LTD	5MG	A077985	001	Apr 23, 2007
	10MG	A077985	002	Apr 23, 2007
HIKMA	5MG	A078129	001	Apr 30, 2008
	10MG	A078129	002	Apr 30, 2008
INVAGEN PHARMS	5MG	A078184	001	Sep 07, 2007
	10MG	A078184	002	Sep 07, 2007
MYLAN PHARMS INC	5MG	A078016	001	Apr 23, 2007
	10MG	A078016	002	Apr 23, 2007
RISING	5MG	A076578	001	Apr 23, 2007
	10MG	A076578	002	Apr 23, 2007
STRIDES PHARMA	5MG	A076062	001	Apr 23, 2007
	5MG	A078616	001	Nov 21, 2008
	10MG	A076062	002	Apr 23, 2007
	10MG	A078616	002	Nov 21, 2008
SUN PHARM INDS INC	5MG	A077359	001	Apr 23, 2007
	10MG	A077359	002	Apr 23, 2007
SUN PHARM INDS LTD	5MG	A078055	001	Apr 23, 2007
	10MG	A078055	002	Apr 23, 2007
SUN PHARM INDUSTRIES	5MG	A077288	001	Apr 23, 2007
	10MG	A077288	002	Apr 23, 2007
SYNTHON PHARMS	5MG	A077540	001	Apr 23, 2007

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ZOLPIDEM TARTRATE

TABLET;ORAL

ZOLPIDEM TARTRATE

	10MG	A077540 002	Apr 23, 2007
WATSON LABS	5MG	A077773 001	Apr 23, 2007
	10MG	A077773 002	Apr 23, 2007
WOCKHARDT	5MG	A078426 001	May 15, 2007
	10MG	A078426 002	May 15, 2007
YUNG SHIN PHARM	5MG	A077990 001	Apr 23, 2007
	10MG	A077990 002	Apr 23, 2007

TABLET;SUBLINGUAL

INTERMEZZO

+	PURDUE PHARMA	1.75MG	N022328 001	Nov 23, 2011
+		3.5MG	N022328 002	Nov 23, 2011

ZOLPIDEM TARTRATE

NORVIUM BIOSCIENCE	5MG	A202657 001	Aug 08, 2016
	10MG	A202657 002	Aug 08, 2016

TABLET, EXTENDED RELEASE;ORAL

ZOLPIDEM TARTRATE

ACTAVIS ELIZABETH	6.25MG	A078179 002	Oct 13, 2010
	12.5MG	A078179 001	Jun 06, 2011
ACTAVIS LABS FL INC	6.25MG	A090153 001	Mar 25, 2013
	12.5MG	A090153 002	Mar 25, 2013
BRECKENRIDGE	6.25MG	A213592 001	Jun 04, 2020
	12.5MG	A213592 002	Jun 04, 2020
ENDO OPERATIONS	6.25MG	A078148 002	Apr 14, 2011
	12.5MG	A078148 001	Dec 03, 2010
SYNTHON PHARMS	6.25MG	A078483 001	Apr 12, 2011
	12.5MG	A078483 002	Jun 06, 2011

TABLET, ORALLY DISINTEGRATING;ORAL

TOVALT ODT

+	BIOVAIL LABS INTL	5MG	N021412 001	Apr 25, 2007
+		10MG	N021412 002	Apr 25, 2007

ZONISAMIDE

CAPSULE;ORAL

ZONEGRAN

+	ADVANZ PHARMA	50MG **	N020789 002	Aug 22, 2003
---	---------------	---------	-------------	--------------

ZONISAMIDE

ANI PHARMS	25MG	A077639 001	Dec 22, 2005
	25MG	A077641 003	Dec 22, 2005
	50MG	A077639 002	Dec 22, 2005
	50MG	A077641 002	Dec 22, 2005
	100MG	A077639 003	Dec 22, 2005
	100MG	A077641 001	Dec 22, 2005
EPIC PHARMA LLC	25MG	A077876 001	Feb 21, 2007
	50MG	A077876 002	Feb 21, 2007
	100MG	A077876 003	Feb 21, 2007
HERITAGE PHARMA AVET	25MG	A077650 001	Apr 20, 2006
	50MG	A077650 002	Apr 20, 2006
	100MG	A077650 003	Apr 20, 2006
NORVIUM BIOSCIENCE	25MG	A077647 001	Dec 22, 2005
	50MG	A077647 002	Dec 22, 2005
	100MG	A077647 003	Dec 22, 2005
RISING	25MG	A077637 001	Dec 22, 2005
	50MG	A077637 002	Dec 22, 2005
	100MG	A077637 003	Dec 22, 2005
ROXANE	25MG	A077648 001	Dec 22, 2005
	50MG	A077648 002	Dec 22, 2005
	100MG	A077648 003	Dec 22, 2005
SUN PHARM INDUSTRIES	25MG	A077635 001	Dec 22, 2005
	50MG	A077635 002	Dec 22, 2005
	100MG	A077635 003	Dec 22, 2005
UPSHER SMITH LABS	25MG	A077644 001	Dec 22, 2005
	50MG	A077644 002	Dec 22, 2005
	100MG	A077644 003	Dec 22, 2005

ORPHAN PRODUCTS DESIGNATIONS AND APPROVALS LIST

The list of Orphan Designations and Approvals is available at:

<http://www.fda.gov/ForIndustry/DevelopingProductsforRareDiseasesConditions/default.htm>

**DRUG PRODUCTS WHICH MUST DEMONSTRATE *IN VIVO* BIOAVAILABILITY
ONLY IF PRODUCT FAILS TO ACHIEVE ADEQUATE DISSOLUTION**

ACETAMINOPHEN;ASPIRIN;BUTALBITAL
CAPSULE OR TABLET; ORAL
160-165MG;160-165MG;50MG
325MG;325MG;50MG

ASPIRIN;CAFFEINE;CARISOPRODOL;
CODEINE PHOSPHATE
TABLET; ORAL
160MG;32MG;200MG;16MG

ACETAMINOPHEN;ASPIRIN;BUTALBITAL;
CAFFEINE
CAPSULE OR TABLET; ORAL
160-165MG;160-165MG;50MG;40MG
325MG;325MG;50MG;40MG

ASPIRIN;CARISOPRODOL
TABLET; ORAL
325MG;200MG

ACETAMINOPHEN;BUTALBITAL
CAPSULE OR TABLET; ORAL
325MG;50MG

ASPIRIN;CARISOPRODOL;
CODEINE PHOSPHATE
TABLET; ORAL
325MG;200MG;16MG

ACETAMINOPHEN;BUTALBITAL;CAFFEINE
CAPSULE OR TABLET; ORAL
325MG;50MG;40MG

ASPIRIN;MEPROBAMATE
TABLET; ORAL
325MG;200MG

AMINOPHYLLINE
TABLET; ORAL
100MG;200MG

ASPIRIN;METHOCARBAMOL
TABLET; ORAL
325MG;400MG

ASPIRIN;BUTALBITAL
CAPSULE OR TABLET; ORAL
325MG;50MG
650MG;50MG

CHLOROTHIAZIDE
TABLET; ORAL
250MG

ASPIRIN;BUTALBITAL;CAFFEINE
CAPSULE OR TABLET; ORAL
325MG;50MG;40MG
650MG;50MG;40MG

HYDROXYZINE HYDROCHLORIDE
TABLET; ORAL
10MG;25MG;
50MG;100MG

ASPIRIN;CAFFEINE;CARISOPRODOL
TABLET; ORAL
160MG;32MG;200MG

PREDNISONE
TABLET; ORAL
1MG;2.5MG;5MG;10MG;
20MG;25MG;50MG

APPENDIX A - PRODUCT NAME INDEX**** A ****

ABACAVIR SULFATE, ABACAVIR SULFATE
ABACAVIR SULFATE AND LAMIVUDINE, ABACAVIR SULFATE
ABACAVIR SULFATE, LAMIVUDINE AND ZIDOVUDINE, ABACAVIR SULFATE
ABELCET, AMPHOTERICIN B
ABILIFY, ARIPIPIRAZOLE
ABILIFY ASIMTUFII, ARIPIPIRAZOLE
ABILIFY MAINTENA KIT, ARIPIPIRAZOLE
ABIRATERONE ACETATE, ABIRATERONE ACETATE
ABLYSINOL, ALCOHOL
ABRAXANE, PACLITAXEL
ABREVA, DOCOSANOL (OTC)
ABSORICA, ISOTRETINOIN
ABSORICA LD, ISOTRETINOIN
ACAMPROSATE CALCIUM, ACAMPROSATE CALCIUM
ACANYA, BENZOYL PEROXIDE
ACARBOSE, ACARBOSE
ACCOLATE, ZAFIRLUKAST
ACCRUFER, FERRIC MALTOL
ACEBUTOLOL HYDROCHLORIDE, ACEBUTOLOL HYDROCHLORIDE
ACETADOTE, ACETYLCYSTEINE
ACETAMINOPHEN, ACETAMINOPHEN (OTC)
ACETAMINOPHEN, ACETAMINOPHEN
ACETAMINOPHEN AND CODEINE PHOSPHATE, ACETAMINOPHEN
ACETAMINOPHEN AND IBUPROFEN, ACETAMINOPHEN (OTC)
ACETAMINOPHEN, ASPIRIN AND CAFFEINE, ACETAMINOPHEN (OTC)
ACETAZOLAMIDE, ACETAZOLAMIDE
ACETAZOLAMIDE SODIUM, ACETAZOLAMIDE SODIUM
ACETIC ACID, ACETIC ACID, GLACIAL
ACETIC ACID 0.25% IN PLASTIC CONTAINER, ACETIC ACID, GLACIAL
ACETYLCYSTEINE, ACETYLCYSTEINE
ACHROMYCIN V, TETRACYCLINE HYDROCHLORIDE
ACIPHEX, RABEPRAZOLE SODIUM
ACITRETIN, ACITRETIN
ACTHAR GEL, CORTICOTROPIN
ACTHAR GEL (AUTOINJECTOR), CORTICOTROPIN
ACTICLATE, DOXYCYCLINE HYCLATE
ACTIGALL, URSODIOL
ACTIVELLA, ESTRADIOL
ACTONEL, RISEDRONATE SODIUM
ACTOPLUS MET, METFORMIN HYDROCHLORIDE
ACTOS, PIOGLITAZONE HYDROCHLORIDE
ACULAR, KETOROLAC TROMETHAMINE
ACULAR LS, KETOROLAC TROMETHAMINE
ACUVAIL, KETOROLAC TROMETHAMINE
ACYCLOVIR, ACYCLOVIR
ACYCLOVIR SODIUM, ACYCLOVIR SODIUM
ACZONE, DAPSONE
ADAPALENE, ADAPALENE (OTC)
ADAPALENE, ADAPALENE
ADAPALENE AND BENZOYL PEROXIDE, ADAPALENE
ADASUVE, LOXAPINE
ADCIRCA, TADALAFIL
ADDERALL XR 10, AMPHETAMINE ASPARTATE
ADDERALL XR 15, AMPHETAMINE ASPARTATE
ADDERALL XR 20, AMPHETAMINE ASPARTATE
ADDERALL XR 25, AMPHETAMINE ASPARTATE
ADDERALL XR 30, AMPHETAMINE ASPARTATE
ADDERALL XR 5, AMPHETAMINE ASPARTATE
ADDYI, FLIBANSERIN
ADEFOVIR DIPIVOXIL, ADEFOVIR DIPIVOXIL
ADEMPAS, RIOCIQUAT
ADENOSINE, ADENOSINE
ADIPEX-P, PHENTERMINE HYDROCHLORIDE
ADLARITY, DONEPEZIL HYDROCHLORIDE
ADRENACLICK, EPINEPHRINE

APPENDIX A - PRODUCT NAME INDEX

** A **

ADRENALIN, EPINEPHRINE
ADREVIEW, IOBENGUANE SULFATE I-123
ADVAIR DISKUS 100/50, FLUTICASONE PROPIONATE
ADVAIR DISKUS 250/50, FLUTICASONE PROPIONATE
ADVAIR DISKUS 500/50, FLUTICASONE PROPIONATE
ADVAIR HFA, FLUTICASONE PROPIONATE
ADVIL, IBUPROFEN (OTC)
ADVIL, IBUPROFEN SODIUM (OTC)
ADVIL ALLERGY AND CONGESTION RELIEF, CHLORPHENIRAMINE MALEATE (OTC)
ADVIL ALLERGY SINUS, CHLORPHENIRAMINE MALEATE (OTC)
ADVIL COLD AND SINUS, IBUPROFEN (OTC)
ADVIL CONGESTION RELIEF, IBUPROFEN (OTC)
ADVIL DUAL ACTION WITH ACETAMINOPHEN, ACETAMINOPHEN (OTC)
ADVIL LIQUI-GELS, IBUPROFEN (OTC)
ADVIL MIGRAINE LIQUI-GELS, IBUPROFEN (OTC)
ADVIL MULTI-SYMPTOM COLD & FLU, CHLORPHENIRAMINE MALEATE (OTC)
ADVIL PM, DIPHENHYDRAMINE CITRATE (OTC)
ADVIL PM, DIPHENHYDRAMINE HYDROCHLORIDE (OTC)
ADZENYS XR-ODT, AMPHETAMINE
AFINITOR, EVEROLIMUS
AFINITOR DISPERZ, EVEROLIMUS
AFIRMELLE, ETHINYL ESTRADIOL
AGAMREE, VAMOROLONE
AGGRASTAT, TIROFIBAN HYDROCHLORIDE
AGRYLIN, ANAGRELIDE HYDROCHLORIDE
AIRDUO RESPICLICK, FLUTICASONE PROPIONATE
AIRSUPRA, ALBUTEROL SULFATE
AK-FLUOR 10%, FLUORESCEIN SODIUM
AK-FLUOR 25%, FLUORESCEIN SODIUM
AKEEGA, ABIRATERONE ACETATE
AKLIEF, TRIFAROTENE
AKOVAZ, EPHEDRINE SULFATE
AKPENTOLATE, CYCLOPENTOLATE HYDROCHLORIDE
AKTEN, LIDOCAINE HYDROCHLORIDE
AKYNZEO, FOSNETUPITANT CHLORIDE HYDROCHLORIDE
AKYNZEO, NETUPITANT
ALA-CORT, HYDROCORTISONE
ALA-SCALP, HYDROCORTISONE
ALAVERT, LORATADINE (OTC)
ALAWAY, KETOTIFEN FUMARATE (OTC)
ALBENDAZOLE, ALBENDAZOLE
ALBUTEROL SULFATE, ALBUTEROL SULFATE
ALBUTEROL SULFATE AND IPRATROPIUM BROMIDE, ALBUTEROL SULFATE
ALCAFTADINE, ALCAFTADINE (OTC)
ALCAINE, PROPARACAINE HYDROCHLORIDE
ALCLOMETASONE DIPROPIONATE, ALCLOMETASONE DIPROPIONATE
ALDACTAZIDE, HYDROCHLOROTHIAZIDE
ALDACTONE, SPIRONOLACTONE
ALECENSA, ALECTINIB HYDROCHLORIDE
ALENDRONATE SODIUM, ALENDRONATE SODIUM
ALEVE, NAPROXEN SODIUM (OTC)
ALEVE PM, DIPHENHYDRAMINE HYDROCHLORIDE (OTC)
ALFENTA, ALFENTANIL HYDROCHLORIDE
ALFENTANIL, ALFENTANIL HYDROCHLORIDE
ALFUZOSIN HYDROCHLORIDE, ALFUZOSIN HYDROCHLORIDE
ALIMTA, PEMETREXED DISODIUM
ALISKIREN HEMIFUMARATE, ALISKIREN HEMIFUMARATE
ALKINDI SPRINKLE, HYDROCORTISONE
ALLEGRA ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
ALLEGRA HIVES, FEXOFENADINE HYDROCHLORIDE (OTC)
ALLEGRA-D 12 HOUR ALLERGY AND CONGESTION, FEXOFENADINE HYDROCHLORIDE (OTC)
ALLEGRA-D 24 HOUR ALLERGY AND CONGESTION, FEXOFENADINE HYDROCHLORIDE (OTC)
ALLI, ORLISTAT (OTC)
ALLOPURINOL, ALLOPURINOL
ALLOPURINOL SODIUM, ALLOPURINOL SODIUM

APPENDIX A - PRODUCT NAME INDEX

** A **

ALLZITAL, ACETAMINOPHEN
ALMOTRIPTAN MALATE, ALMOTRIPTAN MALATE
ALOMIDE, LODOXAMIDE TROMETHAMINE
ALOPRIM, ALLOPURINOL SODIUM
ALOSETRON HYDROCHLORIDE, ALOSETRON HYDROCHLORIDE
ALPHAGAN P, BRIMONIDINE TARTRATE
ALPRAZOLAM, ALPRAZOLAM
ALPROSTADIL, ALPROSTADIL
ALREX, LOTEPREDNOL ETABONATE
ALTACE, RAMIPRIL
ALTAFLUOR BENOX, BENOXINATE HYDROCHLORIDE
ALTAVERA, ETHINYL ESTRADIOL
ALTOPREV, LOVASTATIN
ALTRENO, TRETINOIN
ALUNBRIG, BRIGATINIB
ALVAIZ, ELTROMBOPAG CHOLINE
ALVESCO, CICLESONIDE
ALVIMOPAN, ALVIMOPAN
ALYACEN 1/35, ETHINYL ESTRADIOL
ALYACEN 7/7/7, ETHINYL ESTRADIOL
ALYFTREK, DEUTIVACAFTOR
ALYQ, TADALAFIL
AMABELZ, ESTRADIOL
AMANTADINE HYDROCHLORIDE, AMANTADINE HYDROCHLORIDE
AMARYL, GLIMEPIRIDE
AMBIEN, ZOLPIDEM TARTRATE
AMBIEN CR, ZOLPIDEM TARTRATE
AMBISOME, AMPHOTERICIN B
AMBRISENTAN, AMBRISENTAN
AMCINONIDE, AMCINONIDE
AMELUZ, AMINOLEVULINIC ACID HYDROCHLORIDE
AMICAR, AMINOCAPROIC ACID
AMIDATE, ETOMIDATE
AMIKACIN SULFATE, AMIKACIN SULFATE
AMILORIDE HYDROCHLORIDE, AMILORIDE HYDROCHLORIDE
AMILORIDE HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, AMILORIDE HYDROCHLORIDE
AMINO ACIDS, AMINO ACIDS
AMINOACETIC ACID 1.5% IN PLASTIC CONTAINER, GLYCINE
AMINOCAPROIC ACID, AMINOCAPROIC ACID
AMINOCAPROIC ACID IN PLASTIC CONTAINER, AMINOCAPROIC ACID
AMINOPHYLLINE, AMINOPHYLLINE
AMINOSYN II 10% IN PLASTIC CONTAINER, AMINO ACIDS
AMINOSYN II 15% IN PLASTIC CONTAINER, AMINO ACIDS
AMINOSYN-PF 10%, AMINO ACIDS
AMINOSYN-PF 7%, AMINO ACIDS
AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE
AMITIZA, LUBIPROSTONE
AMITRIPTYLINE HYDROCHLORIDE, AMITRIPTYLINE HYDROCHLORIDE
AMLODIPINE AND OLMESARTAN MEDOXOMIL, AMLODIPINE BESYLATE
AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
AMLODIPINE BESYLATE AND ATORVASTATIN CALCIUM, AMLODIPINE BESYLATE
AMLODIPINE BESYLATE AND BENAZEPRIL HYDROCHLORIDE, AMLODIPINE BESYLATE
AMLODIPINE BESYLATE AND VALSARTAN, AMLODIPINE BESYLATE
AMLODIPINE BESYLATE, VALSARTAN AND HYDROCHLOROTHIAZIDE, AMLODIPINE BESYLATE
AMMONIA N 13, AMMONIA N-13
AMMONIUM CHLORIDE IN PLASTIC CONTAINER, AMMONIUM CHLORIDE
AMMONIUM LACTATE, AMMONIUM LACTATE
AMNESTEEM, ISOTRETINOIN
AMONDYS 45, CASIMERSSEN
AMOXAPINE, AMOXAPINE
AMOXICILLIN, AMOXICILLIN
AMOXICILLIN AND CLAVULANATE POTASSIUM, AMOXICILLIN
AMOXICILLIN PEDIATRIC, AMOXICILLIN
AMOXIL, AMOXICILLIN
AMPHETAMINE SULFATE, AMPHETAMINE SULFATE

APPENDIX A - PRODUCT NAME INDEX

** A **

AMPHOTERICIN B, AMPHOTERICIN B
AMPICILLIN AND SULBACTAM, AMPICILLIN SODIUM
AMPICILLIN SODIUM, AMPICILLIN SODIUM
AMPICILLIN TRIHYDRATE, AMPICILLIN/AMPICILLIN TRIHYDRATE
AMPYRA, DALFAMPRIDINE
AMRINONE LACTATE, INAMRINONE LACTATE
AMRIX, CYCLOBENZAPRINE HYDROCHLORIDE
AMVUTTRA, VUTRISIRAN SODIUM
AMYVID, FLORBETAPIR F-18
AMZEEQ, MINOCYCLINE HYDROCHLORIDE
AN-SULFUR COLLOID, TECHNETIUM TC-99M SULFUR COLLOID KIT
ANAFRANIL, CLOMIPRAMINE HYDROCHLORIDE
ANAGRELIDE HYDROCHLORIDE, ANAGRELIDE HYDROCHLORIDE
ANAPROX DS, NAPROXEN SODIUM
ANASTROZOLE, ANASTROZOLE
ANCOBON, FLUCYTOSINE
ANDROGEL, TESTOSTERONE
ANDROID 25, METHYLTESTOSTERONE
ANECTINE, SUCCINYLCHOLINE CHLORIDE
ANEXSIA 5/325, ACETAMINOPHEN
ANEXSIA 7.5/325, ACETAMINOPHEN
ANGELIQ, DROSPIRENONE
ANGIOMAX, BIVALIRUDIN
ANGIOMAX RTU, BIVALIRUDIN
ANNOVERA, ETHINYL ESTRADIOL
ANORO ELLIPTA, UMECLIDINIUM BROMIDE
ANTARA (MICRONIZED), FENOFIBRATE
ANTHELIOS 20, AVOBENZONE (OTC)
ANTHELIOS 40, AVOBENZONE (OTC)
ANTHELIOS SX, AVOBENZONE (OTC)
ANTIVERT, MECLIZINE HYDROCHLORIDE
ANUSOL HC, HYDROCORTISONE
APHEXDA, MOTIXAFORTIDE ACETATE
APIXABAN, APIXABAN
APLENZIN, BUPROPION HYDROBROMIDE
APOKYN, APOMORPHINE HYDROCHLORIDE
APOMORPHINE HYDROCHLORIDE, APOMORPHINE HYDROCHLORIDE
APONVIE, APREPITANT
APRACLONIDINE HYDROCHLORIDE, APRACLONIDINE HYDROCHLORIDE
APREMILAST, APREMILAST
APREPITANT, APREPITANT
APRETUDE, CABOTEGRAVIR
APRISO, MESALAMINE
APTENSIO XR, METHYLPHENIDATE HYDROCHLORIDE
APTIOM, ESLICARBAZEPINE ACETATE
APTIVUS, TIPRANAVIR
AQNEURSA, LEVACETYLLAUCINE
AQUASOL A, VITAMIN A PALMITATE
ARAKODA, TAFENOQUINE SUCCINATE
ARANELLE, ETHINYL ESTRADIOL
ARAVA, LEFLUNOMIDE
ARAZLO, TAZAROTENE
ARESTIN, MINOCYCLINE HYDROCHLORIDE
ARFORMOTEROL TARTRATE, ARFORMOTEROL TARTRATE
ARGATROBAN, ARGATROBAN
ARGATROBAN IN SODIUM CHLORIDE, ARGATROBAN
ARICEPT, DONEPEZIL HYDROCHLORIDE
ARIDOL KIT, MANNITOL
ARIKAYCE KIT, AMIKACIN SULFATE
ARIMIDEX, ANASTROZOLE
ARIPIPAZOLE, ARIPIPAZOLE
ARISTADA, ARIPIPAZOLE LAUROXIL
ARISTADA INITIO KIT, ARIPIPAZOLE LAUROXIL
ARIXTRA, FONDAPARINUX SODIUM
ARMODAFINIL, ARMODAFINIL

APPENDIX A - PRODUCT NAME INDEX

** A **

ARNUITY ELLIPTA, FLUTICASONE FUROATE
AROMASIN, EXEMESTANE
ARRANON, NELARABINE
ARSENIC TRIOXIDE, ARSENIC TRIOXIDE
ARTESUNATE, ARTESUNATE
ARTHROTEC, DICLOFENAC SODIUM
ASCLERA, POLIDOCANOL
ASCOR, ASCORBIC ACID
ASENAPINE MALEATE, ASENAPINE MALEATE
ASHLYNA, ETHINYL ESTRADIOL
ASMANEX HFA, MOMETASONE FUROATE
ASMANEX TWISTHALER, MOMETASONE FUROATE
ASPIRIN AND DIPYRIDAMOLE, ASPIRIN
ASTAGRAF XL, TACROLIMUS
ASTEPRO ALLERGY, AZELASTINE HYDROCHLORIDE (OTC)
ATACAND, CANDESARTAN CILEXETIL
ATACAND HCT, CANDESARTAN CILEXETIL
ATAZANAVIR SULFATE, ATAZANAVIR SULFATE
ATELVIA, RISEDRONATE SODIUM
ATENOLOL, ATENOLOL
ATENOLOL AND CHLORTHALIDONE, ATENOLOL
ATHENTIA NEXT, LEVONORGESTREL (OTC)
ATIVAN, LORAZEPAM
ATOMOXETINE HYDROCHLORIDE, ATOMOXETINE HYDROCHLORIDE
ATORVALIQ, ATORVASTATIN CALCIUM
ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM
ATOVAQUONE, ATOVAQUONE
ATOVAQUONE AND PROGUANIL HYDROCHLORIDE, ATOVAQUONE
ATRACURIUM BESYLATE, ATRACURIUM BESYLATE
ATRACURIUM BESYLATE PRESERVATIVE FREE, ATRACURIUM BESYLATE
ATRALIN, TRETINOIN
ATROPINE SULFATE, ATROPINE SULFATE
ATROVENT HFA, IPRATROPIUM BROMIDE
ATTRUBY, ACORAMIDIS HYDROCHLORIDE
AUBAGIO, TERIFLUNOMIDE
AUGMENTIN '125', AMOXICILLIN
AUGMENTIN '250', AMOXICILLIN
AUGMENTIN '875', AMOXICILLIN
AUGMENTIN ES-600, AMOXICILLIN
AUGTYRO, REPOTRECTINIB
AURLUMYN, ILOPROST
AUROVELA 1.5/30, ETHINYL ESTRADIOL
AUROVELA 1/20, ETHINYL ESTRADIOL
AUROVELA 24 FE, ETHINYL ESTRADIOL
AUROVELA FE 1.5/30, ETHINYL ESTRADIOL
AUROVELA FE 1/20, ETHINYL ESTRADIOL
AURYXIA, FERRIC CITRATE
AUSTEDO, DEUTETRABENAZINE
AUSTEDO XR, DEUTETRABENAZINE
AUVELITY, BUPROPION HYDROCHLORIDE
AUVI-Q, EPINEPHRINE
AVAGARD, ALCOHOL (OTC)
AVAGE, TAZAROTENE
AVALIDE, HYDROCHLOROTHIAZIDE
AVANAFIL, AVANAFIL
AVAPRO, IRBESARTAN
AVEED, TESTOSTERONE UNDECANOATE
AVIANE-28, ETHINYL ESTRADIOL
AVITA, TRETINOIN
AVODART, DUTASTERIDE
AVYCAZ, AVIBACTAM SODIUM
AXID AR, NIZATIDINE (OTC)
AXTLE, PEMETREXED DIPOTASSIUM
AXUMIN, FLUCICLOVINE F-18
AYUNA, ETHINYL ESTRADIOL

APPENDIX A - PRODUCT NAME INDEX**** A ****

AYVAKIT, AVAPRITINIB
 AZACITIDINE, AZACITIDINE
 AZACTAM, AZTREONAM
 AZASAN, AZATHIOPRINE
 AZASITE, AZITHROMYCIN
 AZATHIOPRINE, AZATHIOPRINE
 AZATHIOPRINE SODIUM, AZATHIOPRINE SODIUM
 AZELAIC ACID, AZELAIC ACID
 AZELASTINE HYDROCHLORIDE, AZELASTINE HYDROCHLORIDE (OTC)
 AZELASTINE HYDROCHLORIDE, AZELASTINE HYDROCHLORIDE
 AZELASTINE HYDROCHLORIDE ALLERGY, AZELASTINE HYDROCHLORIDE (OTC)
 AZELASTINE HYDROCHLORIDE AND FLUTICASONE PROPIONATE, AZELASTINE HYDROCHLORIDE
 AZELASTINE HYDROCHLORIDE CHILDREN'S ALLERGY, AZELASTINE HYDROCHLORIDE (OTC)
 AZELEX, AZELAIC ACID
 AZILECT, RASAGILINE MESYLATE
 AZITHROMYCIN, AZITHROMYCIN
 AZMIRO, TESTOSTERONE CYPIONATE
 AZOPT, BRINZOLAMIDE
 AZOR, AMLODIPINE BESYLATE
 AZSTARYS, DEXMETHYLPHENIDATE HYDROCHLORIDE
 AZTREONAM, AZTREONAM
 AZULFIDINE, SULFASALAZINE
 AZULFIDINE EN-TABS, SULFASALAZINE

**** B ****

BACITRACIN, BACITRACIN
 BACITRACIN ZINC AND POLYMYXIN B SULFATE, BACITRACIN ZINC
 BACITRACIN-NEOMYCIN-POLYMYXIN W/ HYDROCORTISONE ACETATE, BACITRACIN ZINC
 BACLOFEN, BACLOFEN
 BACTERIOSTATIC SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, SODIUM CHLORIDE
 BACTERIOSTATIC WATER FOR INJECTION IN PLASTIC CONTAINER, STERILE WATER FOR INJECTION
 BACTOCILL IN PLASTIC CONTAINER, OXACILLIN SODIUM
 BACTRIM, SULFAMETHOXAZOLE
 BACTRIM DS, SULFAMETHOXAZOLE
 BAFIERTAM, MONOMETHYL FUMARATE
 BAL, DIMERCAPROL
 BALANCED SALT, CALCIUM CHLORIDE
 BALCOLTRA, ETHINYL ESTRADIOL
 BALSALAZIDE DISODIUM, BALSALAZIDE DISODIUM
 BALVERSA, ERDAFITINIB
 BALZIVA-28, ETHINYL ESTRADIOL
 BANZEL, RUFINAMIDE
 BAQSIMI, GLUCAGON
 BARACLUDE, ENTECAVIR
 BARHEMSYS, AMISULPRIDE
 BAXDELA, DELAFLOXACIN MEGLUMINE
 BEIZRAY, DOCETAXEL
 BEKYREE, DESOGESTREL
 BELBUCA, BUPRENORPHINE HYDROCHLORIDE
 BELEODAQ, BELINOSTAT
 BELRAPZO, BENDAMUSTINE HYDROCHLORIDE
 BELSOMRA, SUVOREXANT
 BENAZEPRIL HYDROCHLORIDE, BENAZEPRIL HYDROCHLORIDE
 BENAZEPRIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, BENAZEPRIL HYDROCHLORIDE
 BENDAMUSTINE HYDROCHLORIDE, BENDAMUSTINE HYDROCHLORIDE
 BENDEKA, BENDAMUSTINE HYDROCHLORIDE
 BENICAR, OLMESARTAN MEDOXOMIL
 BENICAR HCT, HYDROCHLOROTHIAZIDE
 BENZAMYCIN, BENZOYL PEROXIDE
 BENZNIDAZOLE, BENZNIDAZOLE
 BENZONATATE, BENZONATATE
 BENZPHETAMINE HYDROCHLORIDE, BENZPHETAMINE HYDROCHLORIDE
 BENZTROPINE MESYLATE, BENZTROPINE MESYLATE
 BEPOTASTINE BESILATE, BEPOTASTINE BESILATE
 BEPREVE, BEPOTASTINE BESILATE

APPENDIX A - PRODUCT NAME INDEX

** B **

BESIVANCE, BESIFLOXACIN HYDROCHLORIDE
BETA-VAL, BETAMETHASONE VALERATE
BETADINE, POVIDONE-IODINE
BETAGAN, LEVOBUNOLOL HYDROCHLORIDE
BETAINE, BETAINE
BETAMETHASONE ACETATE AND BETAMETHASONE SODIUM PHOSPHATE, BETAMETHASONE ACETATE
BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
BETAMETHASONE VALERATE, BETAMETHASONE VALERATE
BETAPACE, SOTALOL HYDROCHLORIDE
BETAPACE AF, SOTALOL HYDROCHLORIDE
BETAXOLOL HYDROCHLORIDE, BETAXOLOL HYDROCHLORIDE
BETHANECHOL CHLORIDE, BETHANECHOL CHLORIDE
BETHKIS, TOBRAMYCIN
BETIMOL, TIMOLOL
BETOPTIC, BETAXOLOL HYDROCHLORIDE
BETOPTIC S, BETAXOLOL HYDROCHLORIDE
BEVESPI AEROSPHERE, FORMOTEROL FUMARATE
BEXAROTENE, BEXAROTENE
BEYAZ, DROSPIRENONE
BICALUTAMIDE, BICALUTAMIDE
BICILLIN C-R, PENICILLIN G BENZATHINE
BICILLIN C-R 900/300, PENICILLIN G BENZATHINE
BICILLIN L-A, PENICILLIN G BENZATHINE
BICNU, CARMUSTINE
BIDIL, HYDRALAZINE HYDROCHLORIDE
BIJUVA, ESTRADIOL
BIKTARVY, BICTEGRAVIR SODIUM
BIMATOPROST, BIMATOPROST
BINOSTO, ALENDRONATE SODIUM
BIORPHEN, PHENYLEPHRINE HYDROCHLORIDE
BIOSCRUB, CHLORHEXIDINE GLUCONATE (OTC)
BISMUTH SUBCITRATE POTASSIUM, METRONIDAZOLE AND TETRACYCLINE HYDROCHLORIDE, BISMUTH
BISMUTH SUBSALICYLATE, METRONIDAZOLE AND TETRACYCLINE HYDROCHLORIDE, BISMUTH SUBSALICYLATE
BISOPROLOL FUMARATE, BISOPROLOL FUMARATE
BISOPROLOL FUMARATE AND HYDROCHLOROTHIAZIDE, BISOPROLOL FUMARATE
BIVALIRUDIN, BIVALIRUDIN
BLEOMYCIN SULFATE, BLEOMYCIN SULFATE
BLISOVI 24 FE, ETHINYL ESTRADIOL
BLISOVI FE 1.5/30, ETHINYL ESTRADIOL
BLISOVI FE 1/20, ETHINYL ESTRADIOL
BLOXIVERZ, NEOSTIGMINE METHYLSULFATE
BLUDIGO, INDIGOTINDISULFONATE SODIUM
BONJESTA, DOXYLAMINE SUCCINATE
BONSITY, TERIPARATIDE
BONTRIL PDM, PHENDIMETRAZINE TARTRATE
BORTEZOMIB, BORTEZOMIB
BOSENTAN, BOSENTAN
BOSULIF, BOSUTINIB MONOHYDRATE
BRAFTOVI, ENCORAFENIB
BRENZAVVY, BEXAGLIFLOZIN
BREO ELLIPTA, FLUTICASONE FUROATE
BRETHINE, TERBUTALINE SULFATE
BREVIBLOC, ESMOLOL HYDROCHLORIDE
BREVIBLOC DOUBLE STRENGTH IN PLASTIC CONTAINER, ESMOLOL HYDROCHLORIDE
BREVIBLOC IN PLASTIC CONTAINER, ESMOLOL HYDROCHLORIDE
BREVITAL SODIUM, METHOHEXITAL SODIUM
BREXAFEMME, IBREXAFUNGERP CITRATE
BREYNA, BUDESONIDE
BREZTRI AEROSPHERE, BUDESONIDE
BRIDION, SUGAMMADEX SODIUM
BRIELLYN, ETHINYL ESTRADIOL
BRILINTA, TICAGRELOR
BRIMONIDINE TARTRATE, BRIMONIDINE TARTRATE (OTC)
BRIMONIDINE TARTRATE, BRIMONIDINE TARTRATE
BRIMONIDINE TARTRATE AND TIMOLOL MALEATE, BRIMONIDINE TARTRATE

APPENDIX A - PRODUCT NAME INDEX

** B **

BRINZOLAMIDE, BRINZOLAMIDE
 BRISDELLE, PAROXETINE MESYLATE
 BRIVIACT, BRIVARACETAM
 BRIXADI, BUPRENORPHINE
 BROMFED-DM, BROMPHENIRAMINE MALEATE
 BROMFENAC SODIUM, BROMFENAC SODIUM
 BROMOCRIPTINE MESYLATE, BROMOCRIPTINE MESYLATE
 BROMPHENIRAMINE MALEATE, PSEUDOEPHEDRINE HYDROCHLORIDE AND DEXTROMETHORPHAN HYDROBROMIDE,
 BROMSITE, BROMFENAC SODIUM
 BRONCHITOL, MANNITOL
 BROVANA, ARFORMOTEROL TARTRATE
 BRUKINSA, ZANUBRUTINIB
 BRYHALI, HALOBETASOL PROPIONATE
 BSS, CALCIUM CHLORIDE
 BSS PLUS, CALCIUM CHLORIDE
 BUDESONIDE, BUDESONIDE (OTC)
 BUDESONIDE, BUDESONIDE
 BUMETANIDE, BUMETANIDE
 BUMEX, BUMETANIDE
 BUPHENYL, SODIUM PHENYL BUTYRATE
 BUPIVACAINE HYDROCHLORIDE, BUPIVACAINE HYDROCHLORIDE
 BUPIVACAINE HYDROCHLORIDE AND EPINEPHRINE, BUPIVACAINE HYDROCHLORIDE
 BUPIVACAINE HYDROCHLORIDE PRESERVATIVE FREE, BUPIVACAINE HYDROCHLORIDE
 BUPIVACAINE LIPOSOME, BUPIVACAINE
 BUPRENORPHINE, BUPRENORPHINE
 BUPRENORPHINE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE
 BUPRENORPHINE HYDROCHLORIDE AND NALOXONE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE
 BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
 BUSPIRONE HYDROCHLORIDE, BUSPIRONE HYDROCHLORIDE
 BUSULFAN, BUSULFAN
 BUSULFEX, BUSULFAN
 BUTALBITAL AND ACETAMINOPHEN, ACETAMINOPHEN
 BUTALBITAL, ACETAMINOPHEN AND CAFFEINE, ACETAMINOPHEN
 BUTALBITAL, ACETAMINOPHEN, CAFFEINE AND CODEINE PHOSPHATE, ACETAMINOPHEN
 BUTALBITAL, ASPIRIN AND CAFFEINE, ASPIRIN
 BUTALBITAL, ASPIRIN, CAFFEINE, AND CODEINE PHOSPHATE, ASPIRIN
 BUTAPAP, ACETAMINOPHEN
 BUTENAFINE HYDROCHLORIDE, BUTENAFINE HYDROCHLORIDE (OTC)
 BUTORPHANOL TARTRATE, BUTORPHANOL TARTRATE
 BUTORPHANOL TARTRATE PRESERVATIVE FREE, BUTORPHANOL TARTRATE
 BUTRANS, BUPRENORPHINE
 BYDUREON BCISE, EXENATIDE SYNTHETIC
 BYETTA, EXENATIDE SYNTHETIC
 BYFAVO, REMIMAZOLAM BESYLATE
 BYLVAY, ODEVIXIBAT
 BYNFEZIA PEN, OCTREOTIDE ACETATE
 BYSTOLIC, NEBIVOLOL HYDROCHLORIDE

** C **

CABAZITAXEL, CABAZITAXEL
 CABENUVA KIT, CABOTEGRAVIR
 CABERGOLINE, CABERGOLINE
 CABOMETYX, CABOZANTINIB S-MALATE
 CABTREO, ADAPALENE
 CADUET, AMLODIPINE BESYLATE
 CAFKIT, CAFFEINE CITRATE
 CAFFEINE CITRATE, CAFFEINE CITRATE
 CALCIPOTRIENE, CALCIPOTRIENE
 CALCIPOTRIENE AND BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
 CALCIPOTRIENE AND BETHAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
 CALCITONIN-SALMON, CALCITONIN SALMON
 CALCITRIOL, CALCITRIOL
 CALCIUM ACETATE, CALCIUM ACETATE
 CALCIUM CHLORIDE 10%, CALCIUM CHLORIDE
 CALCIUM CHLORIDE 10% IN PLASTIC CONTAINER, CALCIUM CHLORIDE

APPENDIX A - PRODUCT NAME INDEX

** C **

CALCIUM GLUCONATE, CALCIUM GLUCONATE
CALCIUM GLUCONATE IN SODIUM CHLORIDE, CALCIUM GLUCONATE
CALDOLOR, IBUPROFEN
CALQUENCE, ACALABRUTINIB
CALQUENCE, ACALABRUTINIB MALEATE
CAMBIA, DICLOFENAC POTASSIUM
CAMCEVI KIT, LEUPROLIDE MESYLATE
CAMILA, NORETHINDRONE
CAMPTOSAR, IRINOTECAN HYDROCHLORIDE
CAMZYOS, MAVACAMTEN
CANASA, MESALAMINE
CANDESARTAN CILEXETIL, CANDESARTAN CILEXETIL
CANDESARTAN CILEXETIL AND HYDROCHLOROTHIAZIDE, CANDESARTAN CILEXETIL
CAPECITABINE, CAPECITABINE
CAPITAL SOLEIL 15, AVOBENZONE (OTC)
CAPLYTA, LUMATEPERONE TOSYLATE
CAPRELSA, VANDETANIB
CAPTOPRIL, CAPTOPRIL
CAPTOPRIL AND HYDROCHLOROTHIAZIDE, CAPTOPRIL
CARAC, FLUOROURACIL
CARAFATE, SUCRALFATE
CARBAGLU, CARGLUMIC ACID
CARBAMAZEPINE, CARBAMAZEPINE
CARBATROL, CARBAMAZEPINE
CARBIDOPA, CARBIDOPA
CARBIDOPA AND LEVODOPA, CARBIDOPA
CARBIDOPA, LEVODOPA AND ENTACAPONE, CARBIDOPA
CARBINOXAMINE MALEATE, CARBINOXAMINE MALEATE
CARBOCAINE, MEPIVACAINE HYDROCHLORIDE
CARBOPLATIN, CARBOPLATIN
CARBOPROST TROMETHAMINE, CARBOPROST TROMETHAMINE
CARDENE IN 0.83% SODIUM CHLORIDE IN PLASTIC CONTAINER, NICARDIPINE HYDROCHLORIDE
CARDENE IN 0.86% SODIUM CHLORIDE IN PLASTIC CONTAINER, NICARDIPINE HYDROCHLORIDE
CARDENE IN 4.8% DEXTROSE IN PLASTIC CONTAINER, NICARDIPINE HYDROCHLORIDE
CARDIOGEN-82, RUBIDIUM CHLORIDE RB-82
CARDIOLITE, TECHNETIUM TC-99M SESTAMIBI KIT
CARDIOPLEGIC IN PLASTIC CONTAINER, CALCIUM CHLORIDE
CARDIZEM, DILTIAZEM HYDROCHLORIDE
CARDIZEM CD, DILTIAZEM HYDROCHLORIDE
CARDIZEM LA, DILTIAZEM HYDROCHLORIDE
CARDURA, DOXAZOSIN MESYLATE
CARDURA XL, DOXAZOSIN MESYLATE
CARFILZOMIB, CARFILZOMIB
CARGLUMIC ACID, CARGLUMIC ACID
CARISOPRODOL, CARISOPRODOL
CARMUSTINE, CARMUSTINE
CARNITOR, LEVOCARNITINE
CARNITOR SF, LEVOCARNITINE
CAROSPIR, SPIRONOLACTONE
CARTEOLOL HYDROCHLORIDE, CARTEOLOL HYDROCHLORIDE
CARTIA XT, DILTIAZEM HYDROCHLORIDE
CARVEDILOL, CARVEDILOL
CARVEDILOL PHOSPHATE, CARVEDILOL PHOSPHATE
CASODEX, BICALUTAMIDE
CASPOFUNGIN ACETATE, CASPOFUNGIN ACETATE
CASPORYN HC, HYDROCORTISONE
CATAFLAM, DICLOFENAC POTASSIUM
CATAPRES-TTS-1, CLONIDINE
CATAPRES-TTS-2, CLONIDINE
CATAPRES-TTS-3, CLONIDINE
CAVERJECT, ALPROSTADIL
CAVERJECT IMPULSE, ALPROSTADIL
CAYSTON, AZTREONAM
CEFAFLOR, CEFAFLOR
CEFADROXIL, CEFADROXIL/CEFADROXIL HEMIHYDRATE

APPENDIX A - PRODUCT NAME INDEX

** C **

CEFAZOLIN AND DEXTROSE, CEFAZOLIN SODIUM
CEFAZOLIN IN DEXTROSE, CEFAZOLIN SODIUM
CEFAZOLIN SODIUM, CEFAZOLIN SODIUM
CEFDINIR, CEFDINIR
CEFEPIME AND DEXTROSE IN DUPLEX CONTAINER, CEFEPIME HYDROCHLORIDE
CEFEPIME HYDROCHLORIDE, CEFEPIME HYDROCHLORIDE
CEFEPIME HYDROCHLORIDE IN PLASTIC CONTAINER, CEFEPIME HYDROCHLORIDE
CEFEPIME IN PLASTIC CONTAINER, CEFEPIME HYDROCHLORIDE
CEFIXIME, CEFIXIME
CEFOTAN, CEFOTETAN DISODIUM
CEFOTETAN, CEFOTETAN DISODIUM
CEFOXITIN, CEFOXITIN SODIUM
CEFOXITIN AND DEXTROSE IN DUPLEX CONTAINER, CEFOXITIN SODIUM
CEFOXITIN IN PLASTIC CONTAINER, CEFOXITIN SODIUM
CEFPODOXIME PROXETIL, CEFPODOXIME PROXETIL
CEFPROZIL, CEFPROZIL
CEFTAZIDIME, CEFTAZIDIME
CEFTRIAXONE, CEFTRIAXONE SODIUM
CEFTRIAXONE AND DEXTROSE IN DUPLEX CONTAINER, CEFTRIAXONE SODIUM
CEFTRIAXONE IN PLASTIC CONTAINER, CEFTRIAXONE SODIUM
CEFTRIAXONE SODIUM, CEFTRIAXONE SODIUM
CEFUROXIME AXETIL, CEFUROXIME AXETIL
CEFUROXIME SODIUM, CEFUROXIME SODIUM
CELEBEX, CELECOXIB
CELECOXIB, CELECOXIB
CELESTONE SOLUSPAN, BETAMETHASONE ACETATE
CELEXA, CITALOPRAM HYDROBROMIDE
CELLCEPT, MYCOPHENOLATE MOFETIL
CELLCEPT, MYCOPHENOLATE MOFETIL HYDROCHLORIDE
CELONTIN, METHSUXIMIDE
CENTANY, MUPIROCIN
CEPHALEXIN, CEPHALEXIN
CEQUA, CYCLOSPORINE
CERDELGA, ELIGLUSTAT TARTRATE
CEREBYX, FOSPHENYTOIN SODIUM
CERETEC, TECHNETIUM TC-99M EXAMETAZIME KIT
CERIANNA, FLUOROESTRADIOL F-18
CERUBIDINE, DAUNORUBICIN HYDROCHLORIDE
CERVIDIL, DINOPROSTONE
CETIRIZINE HYDROCHLORIDE, CETIRIZINE HYDROCHLORIDE (OTC)
CETIRIZINE HYDROCHLORIDE, CETIRIZINE HYDROCHLORIDE
CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
CETIRIZINE HYDROCHLORIDE AND PSEUDOEPHEDRINE HYDROCHLORIDE, CETIRIZINE HYDROCHLORIDE
CETIRIZINE HYDROCHLORIDE HIVES, CETIRIZINE HYDROCHLORIDE (OTC)
CETIRIZINE HYDROCHLORIDE HIVES RELIEF, CETIRIZINE HYDROCHLORIDE (OTC)
CETRAXAL, CIPROFLOXACIN HYDROCHLORIDE
CETRORELIX ACETATE, CETRORELIX ACETATE
CETROTIDE, CETRORELIX ACETATE
CEVIMELINE HYDROCHLORIDE, CEVIMELINE HYDROCHLORIDE
CHABELINA FE, ETHINYL ESTRADIOL
CHEMET, SUCCIMER
CHENODIOL, CHENODIOL
CHG SCRUB, CHLORHEXIDINE GLUCONATE (OTC)
CHILDREN'S ADVIL, IBUPROFEN (OTC)
CHILDREN'S ADVIL ALLERGY SINUS, CHLORPHENIRAMINE MALEATE (OTC)
CHILDREN'S ADVIL COLD, IBUPROFEN (OTC)
CHILDREN'S ADVIL-FLAVORED, IBUPROFEN (OTC)
CHILDREN'S ALAWAY, KETOTIFEN FUMARATE (OTC)
CHILDREN'S ALLEGRA ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
CHILDREN'S ASTEPRO ALLERGY, AZELASTINE HYDROCHLORIDE (OTC)
CHILDREN'S CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
CHILDREN'S CETIRIZINE HYDROCHLORIDE HIVES RELIEF, CETIRIZINE HYDROCHLORIDE (OTC)
CHILDREN'S CLARITIN, LORATADINE (OTC)
CHILDREN'S FEXOFENADINE HYDROCHLORIDE ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
CHILDREN'S FEXOFENADINE HYDROCHLORIDE HIVES, FEXOFENADINE HYDROCHLORIDE (OTC)

APPENDIX A - PRODUCT NAME INDEX

** C **

CHILDREN'S IBUPROFEN, IBUPROFEN (OTC)
 CHILDREN'S MOTRIN, IBUPROFEN (OTC)
 CHILDREN'S MOTRIN COLD, IBUPROFEN (OTC)
 CHILDREN'S ZYRTEC ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 CHILDREN'S ZYRTEC HIVES, CETIRIZINE HYDROCHLORIDE (OTC)
 CHIRHOSTIM, SECRETIN SYNTHETIC HUMAN
 CHLORAPREP ONE-STEP, CHLORHEXIDINE GLUCONATE (OTC)
 CHLORAPREP ONE-STEP FREPP, CHLORHEXIDINE GLUCONATE (OTC)
 CHLORAPREP ONE-STEP SEPP, CHLORHEXIDINE GLUCONATE (OTC)
 CHLORAPREP SINGLE SWABSTICK, CHLORHEXIDINE GLUCONATE (OTC)
 CHLORAPREP TRIPLE SWABSTICK, CHLORHEXIDINE GLUCONATE (OTC)
 CHLORAPREP WITH TINT, CHLORHEXIDINE GLUCONATE (OTC)
 CHLORDIAZEPOXIDE AND AMITRIPTYLINE HYDROCHLORIDE, AMITRIPTYLINE HYDROCHLORIDE
 CHLORDIAZEPOXIDE HYDROCHLORIDE, CHLORDIAZEPOXIDE HYDROCHLORIDE
 CHLORDIAZEPOXIDE HYDROCHLORIDE AND CLIDINIUM BROMIDE, CHLORDIAZEPOXIDE HYDROCHLORIDE
 CHLORHEXIDINE GLUCONATE, CHLORHEXIDINE GLUCONATE (OTC)
 CHLORHEXIDINE GLUCONATE, CHLORHEXIDINE GLUCONATE
 CHLOROPROCAINE HYDROCHLORIDE, CHLOROPROCAINE HYDROCHLORIDE
 CHLOROQUINE PHOSPHATE, CHLOROQUINE PHOSPHATE
 CHLOROTHIAZIDE SODIUM, CHLOROTHIAZIDE SODIUM
 CHLORPHENIRAMINE MALEATE, CHLORPHENIRAMINE MALEATE (OTC)
 CHLORPROMAZINE HYDROCHLORIDE, CHLORPROMAZINE HYDROCHLORIDE
 CHLORTHALIDONE, CHLORTHALIDONE
 CHLORZOXAZONE, CHLORZOXAZONE
 CHOLBAM, CHOLIC ACID
 CHOLESTYRAMINE, CHOLESTYRAMINE
 CHOLESTYRAMINE LIGHT, CHOLESTYRAMINE
 CHOLETEC, TECHNETIUM TC-99M MEBROFENIN KIT
 CHOLINE C-11, CHOLINE C-11
 CHROMIC CHLORIDE, CHROMIC CHLORIDE
 CHROMIC CHLORIDE IN PLASTIC CONTAINER, CHROMIC CHLORIDE
 CIALIS, TADALAFIL
 CIBINQO, ABROCITINIB
 CICLOPIROX, CICLOPIROX
 CIDA-STAT, CHLORHEXIDINE GLUCONATE (OTC)
 CIDOFOVIR, CIDOFOVIR
 CILOSTAZOL, CILOSTAZOL
 CILOXAN, CIPROFLOXACIN HYDROCHLORIDE
 CIMDUO, LAMIVUDINE
 CIMETIDINE, CIMETIDINE (OTC)
 CIMETIDINE, CIMETIDINE
 CIMETIDINE HYDROCHLORIDE, CIMETIDINE HYDROCHLORIDE
 CINACALCET HYDROCHLORIDE, CINACALCET HYDROCHLORIDE
 CINVANTI, APREPITANT
 CIPRO, CIPROFLOXACIN
 CIPRO, CIPROFLOXACIN HYDROCHLORIDE
 CIPRO HC, CIPROFLOXACIN HYDROCHLORIDE
 CIPRODEX, CIPROFLOXACIN
 CIPROFLOXACIN, CIPROFLOXACIN
 CIPROFLOXACIN AND DEXAMETHASONE, CIPROFLOXACIN
 CIPROFLOXACIN EXTENDED RELEASE, CIPROFLOXACIN
 CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE
 CIPROFLOXACIN IN DEXTROSE 5% IN PLASTIC CONTAINER, CIPROFLOXACIN
 CIPROFLOXACIN; DEXAMETHASONE, CIPROFLOXACIN
 CIS-MDP, TECHNETIUM TC-99M MEDRONATE KIT
 CIS-PYRO, TECHNETIUM TC-99M PYROPHOSPHATE KIT
 CISATRACURIUM BESYLATE, CISATRACURIUM BESYLATE
 CISATRACURIUM BESYLATE PRESERVATIVE FREE, CISATRACURIUM BESYLATE
 CISPLATIN, CISPLATIN
 CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE
 CITANEST FORTE DENTAL, EPINEPHRINE BITARTRATE
 CLADRIBINE, CLADRIBINE
 CLARAVIS, ISOTRETINOIN
 CLARINEX, DESLORATADINE
 CLARINEX-D 12 HOUR, DESLORATADINE

APPENDIX A - PRODUCT NAME INDEX

** C **

CLARISCAN, GADOTERATE MEGLUMINE
 CLARITHROMYCIN, CLARITHROMYCIN
 CLARITIN, LORATADINE (OTC)
 CLARITIN HIVES RELIEF, LORATADINE (OTC)
 CLARITIN HIVES RELIEF REDITAB, LORATADINE (OTC)
 CLARITIN REDITABS, LORATADINE (OTC)
 CLARITIN-D, LORATADINE (OTC)
 CLARITIN-D 24 HOUR, LORATADINE (OTC)
 CLEMASTINE FUMARATE, CLEMASTINE FUMARATE
 CLENPIQ, CITRIC ACID
 CLEOCIN, CLINDAMYCIN PALMITATE HYDROCHLORIDE
 CLEOCIN, CLINDAMYCIN PHOSPHATE
 CLEOCIN HYDROCHLORIDE, CLINDAMYCIN HYDROCHLORIDE
 CLEOCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
 CLEOCIN T, CLINDAMYCIN PHOSPHATE
 CLEVIPREX, CLEVIDIPINE
 CLIMARA, ESTRADIOL
 CLIMARA PRO, ESTRADIOL
 CLINDA-DERM, CLINDAMYCIN PHOSPHATE
 CLINDAGEL, CLINDAMYCIN PHOSPHATE
 CLINDAMYCIN HYDROCHLORIDE, CLINDAMYCIN HYDROCHLORIDE
 CLINDAMYCIN PALMITATE HYDROCHLORIDE, CLINDAMYCIN PALMITATE HYDROCHLORIDE
 CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
 CLINDAMYCIN PHOSPHATE AND BENZOYL PEROXIDE, BENZOYL PEROXIDE
 CLINDAMYCIN PHOSPHATE AND TRETINOIN, CLINDAMYCIN PHOSPHATE
 CLINDAMYCIN PHOSPHATE IN 0.9% SODIUM CHLORIDE, CLINDAMYCIN PHOSPHATE
 CLINDAMYCIN PHOSPHATE IN 5% DEXTROSE IN PLASTIC CONTAINER, CLINDAMYCIN PHOSPHATE
 CLINDAMYCIN PHOSPHATE IN DEXTROSE 5%, CLINDAMYCIN PHOSPHATE
 CLINDESSE, CLINDAMYCIN PHOSPHATE
 CLINDETS, CLINDAMYCIN PHOSPHATE
 CLINIMIX 2.75/10 SULFITE FREE IN DEXTROSE 10% IN PLASTIC CONTAINER, AMINO ACIDS
 CLINIMIX 2.75/25 SULFITE FREE IN DEXTROSE 25% IN PLASTIC CONTAINER, AMINO ACIDS
 CLINIMIX 2.75/5 SULFITE FREE IN DEXTROSE 5% IN PLASTIC CONTAINER, AMINO ACIDS
 CLINIMIX 4.25/10 SULFITE FREE IN DEXTROSE 10% IN PLASTIC CONTAINER, AMINO ACIDS
 CLINIMIX 4.25/20 SULFITE FREE IN DEXTROSE 20% IN PLASTIC CONTAINER, AMINO ACIDS
 CLINIMIX 4.25/25 SULFITE FREE IN DEXTROSE 25% IN PLASTIC CONTAINER, AMINO ACIDS
 CLINIMIX 4.25/5 SULFITE FREE IN DEXTROSE 5% IN PLASTIC CONTAINER, AMINO ACIDS
 CLINIMIX 5/10 SULFITE FREE IN DEXTROSE 10% IN PLASTIC CONTAINER, AMINO ACIDS
 CLINIMIX 5/15 SULFITE FREE IN DEXTROSE 15% IN PLASTIC CONTAINER, AMINO ACIDS
 CLINIMIX 5/20 SULFITE FREE IN DEXTROSE 20% IN PLASTIC CONTAINER, AMINO ACIDS
 CLINIMIX 5/25 SULFITE FREE IN DEXTROSE 25% IN PLASTIC CONTAINER, AMINO ACIDS
 CLINIMIX 5/35 SULFITE FREE IN DEXTROSE 35% IN PLASTIC CONTAINER, AMINO ACIDS
 CLINIMIX 8/14 SULFITE FREE IN DEXTROSE 14% IN PLASTIC CONTAINER, AMINO ACIDS
 CLINIMIX E 2.75/10 SULFITE FREE W/ ELECT IN DEXTROSE 10% W/ CALCIUM IN PLASTIC CONTAINER,
 CLINIMIX E 2.75/25 SULFITE FREE W/ ELECT IN DEXTROSE 25% W/ CALCIUM IN PLASTIC CONTAINER,
 CLINIMIX E 2.75/5 SULFITE FREE W/ ELECT IN DEXTROSE 5% W/ CALCIUM IN PLASTIC CONTAINER,
 CLINIMIX E 4.25/10 SULFITE FREE W/ ELECT IN DEXTROSE 10% W/ CALCIUM IN PLASTIC CONTAINER,
 CLINIMIX E 4.25/20 SULFITE FREE W/ ELECT IN DEXTROSE 20% W/ CALCIUM IN PLASTIC CONTAINER,
 CLINIMIX E 4.25/25 SULFITE FREE W/ ELECT IN DEXTROSE 25% W/ CALCIUM IN PLASTIC CONTAINER,
 CLINIMIX E 4.25/5 SULFITE FREE W/ ELECT IN DEXTROSE 5% W/ CALCIUM IN PLASTIC CONTAINER,
 CLINIMIX E 5/10 SULFITE FREE W/ ELECT IN DEXTROSE 10% W/ CALCIUM IN PLASTIC CONTAINER,
 CLINIMIX E 5/15 SULFITE FREE W/ ELECT IN DEXTROSE 15% W/ CALCIUM IN PLASTIC CONTAINER,
 CLINIMIX E 5/20 SULFITE FREE W/ ELECT IN 20% DEXTROSE W/ CALCIUM IN PLASTIC CONTAINER,
 CLINIMIX E 5/25 SULFITE FREE W/ ELECT IN DEXTROSE 25% W/ CALCIUM IN PLASTIC CONTAINER,
 CLINIMIX E 5/35 SULFITE FREE W/ ELECT IN DEXTROSE 35% W/ CALCIUM IN PLASTIC CONTAINER,
 CLINISOL 15% SULFITE FREE IN PLASTIC CONTAINER, AMINO ACIDS
 CLINOLIPID 20%, OLIVE OIL
 CLOBAZAM, CLOBAZAM
 CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
 CLOBETASOL PROPIONATE (EMOLLIENT), CLOBETASOL PROPIONATE
 CLOBEX, CLOBETASOL PROPIONATE
 CLOCORTOLONE PIVALATE, CLOCORTOLONE PIVALATE
 CLODERM, CLOCORTOLONE PIVALATE
 CLOFARABINE, CLOFARABINE
 CLOLAR, CLOFARABINE

APPENDIX A - PRODUCT NAME INDEX

** C **

CLOMIPHENE CITRATE, CLOMIPHENE CITRATE
CLOMIPRAMINE HYDROCHLORIDE, CLOMIPRAMINE HYDROCHLORIDE
CLONAZEPAM, CLONAZEPAM
CLONIDINE, CLONIDINE
CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
CLOPIDOGREL BISULFATE, CLOPIDOGREL BISULFATE
CLORAZEPATE DIPOTASSIUM, CLORAZEPATE DIPOTASSIUM
CLOTRIMAZOLE, CLOTRIMAZOLE (OTC)
CLOTRIMAZOLE, CLOTRIMAZOLE
CLOTRIMAZOLE AND BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
CLOZAPINE, CLOZAPINE
CLOZARIL, CLOZAPINE
COARTEM, ARTEMETHER
COBENFY, TROSPIMUM CHLORIDE
CODEINE SULFATE, CODEINE SULFATE
COL-PROBENECID, COLCHICINE
COLAZAL, BALSALAZIDE DISODIUM
COLCHICINE, COLCHICINE
COLESEVELAM HYDROCHLORIDE, COLESEVELAM HYDROCHLORIDE
COLESTID, COLESTIPOL HYDROCHLORIDE
COLESTIPOL HYDROCHLORIDE, COLESTIPOL HYDROCHLORIDE
COLISTIMETHATE SODIUM, COLISTIMETHATE SODIUM
COLOCORT, HYDROCORTISONE
COLY-MYCIN M, COLISTIMETHATE SODIUM
COLY-MYCIN S, COLISTIN SULFATE
COMBIGAN, BRIMONIDINE TARTRATE
COMBIPATCH, ESTRADIOL
COMBIVENT RESPIMAT, ALBUTEROL SULFATE
COMBOGESIC, ACETAMINOPHEN
COMBOGESIC IV, ACETAMINOPHEN
COMETRIQ, CABOZANTINIB S-MALATE
COMPLERA, EMTRICITABINE
COMPRO, PROCHLORPERAZINE
COMTAN, ENTACAPONE
CONCERTA, METHYLPHENIDATE HYDROCHLORIDE
CONDYLOX, PODOFILOX
CONRAY, IOTHALAMATE MEGLUMINE
CONTRAVE, BUPROPION HYDROCHLORIDE
CONZIP, TRAMADOL HYDROCHLORIDE
COPAXONE, GLATIRAMER ACETATE
COPIKTRA, DUVELISIB
CORDRAN, FLURANDRENOLIDE
COREG, CARVEDILOL
CORLANOR, IVABRADINE
CORLANOR, IVABRADINE HYDROCHLORIDE
CORPHEDRA, EPHEDRINE SULFATE
CORTEF, HYDROCORTISONE
CORTENEMA, HYDROCORTISONE
CORTIFOAM, HYDROCORTISONE ACETATE
CORTROSYN, COSYNTROPIN
CORVERT, IBUTILIDE FUMARATE
COSELA, TRILACICLIB DIHYDROCHLORIDE
COSOPT, DORZOLAMIDE HYDROCHLORIDE
COSOPT PF, DORZOLAMIDE HYDROCHLORIDE
COSYNTROPIN, COSYNTROPIN
COTELLIC, COBIMETINIB FUMARATE
COTEMPLA XR-ODT, METHYLPHENIDATE
COXANTO, OXAPROZIN
COZAAR, LOSARTAN POTASSIUM
CRENESSITY, CRINECERFONT
CRESEMBA, ISAVUCONAZONIUM SULFATE
CRESTOR, ROSUVASTATIN CALCIUM
CREXONT, CARBIDOPA
CRINONE, PROGESTERONE
CROMOLYN SODIUM, CROMOLYN SODIUM (OTC)

APPENDIX A - PRODUCT NAME INDEX

** C **

CROMOLYN SODIUM, CROMOLYN SODIUM
 CROTAN, CROTAMITON
 CRYSELLE, ETHINYL ESTRADIOL
 CUPRIC CHLORIDE, CUPRIC CHLORIDE
 CUPRIC CHLORIDE IN PLASTIC CONTAINER, CUPRIC CHLORIDE
 CUPRIC SULFATE, CUPRIC SULFATE
 CUPRIMINE, PENICILLAMINE
 CUVPOSA, GLYCOPYRROLATE
 CUVRIOR, TRIENTINE TETRAHYDROCHLORIDE
 CYANOCOBALAMIN, CYANOCOBALAMIN
 CYANOKIT, HYDROXOCOBALAMIN
 CYCLESSA, DESOGESTREL
 CYCLOBENZAPRINE HYDROCHLORIDE, CYCLOBENZAPRINE HYDROCHLORIDE
 CYCLOGYL, CYCLOPENTOLATE HYDROCHLORIDE
 CYCLOMYDRIL, CYCLOPENTOLATE HYDROCHLORIDE
 CYCLOPHOSPHAMIDE, CYCLOPHOSPHAMIDE
 CYCLOSET, BROMOCRIPTINE MESYLATE
 CYCLOSPORINE, CYCLOSPORINE
 CYKLOKAPRON, TRANEXAMIC ACID
 CYMBALTA, DULOXETINE HYDROCHLORIDE
 CYONANZ, ETHINYL ESTRADIOL
 CYPROHEPTADINE HYDROCHLORIDE, CYPROHEPTADINE HYDROCHLORIDE
 CYSTADANE, BETAINE
 CYSTADROPS, CYSTEAMINE HYDROCHLORIDE
 CYSTAGON, CYSTEAMINE BITARTRATE
 CYSTARAN, CYSTEAMINE HYDROCHLORIDE
 CYSTO-CONRAY II, IOTHALAMATE MEGLUMINE
 CYSTOGRAFIN, DIATRIZOATE MEGLUMINE
 CYSTOGRAFIN DILUTE, DIATRIZOATE MEGLUMINE
 CYSVIEW KIT, HEXAMINOLEVULINATE HYDROCHLORIDE
 CYTALUX, PAFOLACIANINE SODIUM
 CYTARABINE, CYTARABINE
 CYTOMEL, LIOTHYRONINE SODIUM
 CYTOTEC, MISOPROSTOL
 CYTOXAN, CYCLOPHOSPHAMIDE

** D **

DABIGATRAN ETEXILATE MESYLATE, DABIGATRAN ETEXILATE MESYLATE
 DACARBAZINE, DACARBAZINE
 DACTINOMYCIN, DACTINOMYCIN
 DALFAMPRIDINE, DALFAMPRIDINE
 DALIRESP, ROFLUMILAST
 DALVANCE, DALBAVANCIN HYDROCHLORIDE
 DANAZOL, DANAZOL
 DANTRIUM, DANTROLENE SODIUM
 DANTROLENE SODIUM, DANTROLENE SODIUM
 DANZITEN, NILOTINIB TARTRATE
 DAPSONE, DAPSONE
 DAPTOMYCIN, DAPTOMYCIN
 DAPTOMYCIN IN 0.9% SODIUM CHLORIDE, DAPTOMYCIN
 DARAPRIM, PYRIMETHAMINE
 DARIFENACIN, DARIFENACIN HYDROBROMIDE
 DARIFENACIN HYDROBROMIDE, DARIFENACIN HYDROBROMIDE
 DARUNAVIR, DARUNAVIR
 DASATINIB, DASATINIB
 DASETTA 1/35, ETHINYL ESTRADIOL
 DASETTA 7/7/7, ETHINYL ESTRADIOL
 DATSCAN, IOFLUPANE I-123
 DAUNORUBICIN HYDROCHLORIDE, DAUNORUBICIN HYDROCHLORIDE
 DAURISMO, GLASDEGIB MALEATE
 DAYBUE, TROFINETIDE
 DAYPRO, OXAPROZIN
 DAYSEE, ETHINYL ESTRADIOL
 DAYTRANA, METHYLPHENIDATE
 DAYVIGO, LEMBOREXANT

APPENDIX A - PRODUCT NAME INDEX

** D **

DDAVP, DESMOPRESSIN ACETATE
DECITABINE, DECITABINE
DEFENCATH, HEPARIN SODIUM
DEFERASIROX, DEFERASIROX
DEFERIPRONE, DEFERIPRONE
DEFEROXAMINE MESYLATE, DEFEROXAMINE MESYLATE
DEFINITY, PERFLUTREN
DEFINITY RT, PERFLUTREN
DEFITELIO, DEFIBROTIDE SODIUM
DEFLAZACORT, DEFLAZACORT
DELESTROGEN, ESTRADIOL VALERATE
DELFLEX W/ DEXTROSE 1.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
DELFLEX W/ DEXTROSE 1.5% LOW MAGNESIUM IN PLASTIC CONTAINER, CALCIUM CHLORIDE
DELFLEX W/ DEXTROSE 1.5% LOW MAGNESIUM LOW CALCIUM IN PLASTIC CONTAINER, CALCIUM CHLORIDE
DELFLEX W/ DEXTROSE 2.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
DELFLEX W/ DEXTROSE 2.5% LOW MAGNESIUM IN PLASTIC CONTAINER, CALCIUM CHLORIDE
DELFLEX W/ DEXTROSE 2.5% LOW MAGNESIUM LOW CALCIUM IN PLASTIC CONTAINER, CALCIUM CHLORIDE
DELFLEX W/ DEXTROSE 4.25% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
DELFLEX W/ DEXTROSE 4.25% LOW MAGNESIUM IN PLASTIC CONTAINER, CALCIUM CHLORIDE
DELFLEX W/ DEXTROSE 4.25% LOW MAGNESIUM LOW CALCIUM IN PLASTIC CONTAINER, CALCIUM CHLORIDE
DELSTRIGO, DORAVIRINE
DELSYM, DEXTROMETHORPHAN POLISTIREX (OTC)
DEMECLOCYCLINE HYDROCHLORIDE, DEMECLOCYCLINE HYDROCHLORIDE
DEMEROL, MEPERIDINE HYDROCHLORIDE
DEMSER, METYROSINE
DENA VIR, PENCICLOVIR
DEOXYCHOLIC ACID, DEOXYCHOLIC ACID
DEPAKOTE, DIVALPROEX SODIUM
DEPAKOTE ER, DIVALPROEX SODIUM
DEPEN, PENICILLAMINE
DEPO-ESTRADIOL, ESTRADIOL CYPIONATE
DEPO-MEDROL, METHYLPREDNISOLONE ACETATE
DEPO-PROVERA, MEDROXYPROGESTERONE ACETATE
DEPO-SUBQ PROVERA 104, MEDROXYPROGESTERONE ACETATE
DEPO-TESTOSTERONE, TESTOSTERONE CYPIONATE
DERMA-SMOOTH/FS, FLUOCINOLONE ACETONIDE
DERMABET, BETAMETHASONE VALERATE
DERMOTIC, FLUOCINOLONE ACETONIDE
DESCOVY, EMTRICITABINE
DESFERAL, DEFEROXAMINE MESYLATE
DESFLURANE, DESFLURANE
DESIPRAMINE HYDROCHLORIDE, DESIPRAMINE HYDROCHLORIDE
DESLORATADINE, DESLORATADINE
DESLORATADINE AND PSEUDOEPHEDRINE SULFATE 24 HOUR, DESLORATADINE
DESMOPRESSIN ACETATE, DESMOPRESSIN ACETATE
DESMOPRESSIN ACETATE (NEEDS NO REFRIGERATION), DESMOPRESSIN ACETATE
DESOGESTREL AND ETHINYL ESTRADIOL, DESOGESTREL
DESONIDE, DESONIDE
DESOWEN, DESONIDE
DESOXIMETASONE, DESOXIMETASONE
DESOXYN, METHAMPHETAMINE HYDROCHLORIDE
DESVENLAFAXINE, DESVENLAFAXINE
DESVENLAFAXINE SUCCINATE, DESVENLAFAXINE SUCCINATE
DETECTNET, COPPER CU-64 DOTATATE
DETROL, TOLTERODINE TARTRATE
DETROL LA, TOLTERODINE TARTRATE
DEUTETRABENAZINE, DEUTETRABENAZINE
DEXAMETHASONE, DEXAMETHASONE
DEXAMETHASONE INTENSOL, DEXAMETHASONE
DEXAMETHASONE SODIUM PHOSPHATE, DEXAMETHASONE SODIUM PHOSPHATE
DEXAMETHASONE SODIUM PHOSPHATE PRESERVATIVE FREE, DEXAMETHASONE SODIUM PHOSPHATE
DEXASPORIN, DEXAMETHASONE
DEXEDRINE SPANSULE, DEXTROAMPHETAMINE SULFATE
DEXILANT, DEXLANSOPRAZOLE
DEXLANSOPRAZOLE, DEXLANSOPRAZOLE

APPENDIX A - PRODUCT NAME INDEX

** D **

DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE
 DEXMETHYLPHENIDATE HYDROCHLORIDE, DEXMETHYLPHENIDATE HYDROCHLORIDE
 DEXRAZOXANE HYDROCHLORIDE, DEXRAZOXANE HYDROCHLORIDE
 DEXTENZA, DEXAMETHASONE
 DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE
 DEXTROAMPHETAMINE SULFATE, DEXTROAMPHETAMINE SULFATE
 DEXTROMETHORPHAN HYDROBROMIDE AND QUINIDINE SULFATE, DEXTROMETHORPHAN HYDROBROMIDE
 DEXTROMETHORPHAN POLISTIREX, DEXTROMETHORPHAN POLISTIREX (OTC)
 DEXTROSE 10% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 10% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 10% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 10% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 10% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 10% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 2.5% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 2.5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 2.5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 2.5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 2.5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 2.5% IN HALF-STRENGTH LACTATED RINGER'S IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 DEXTROSE 20% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 25%, DEXTROSE
 DEXTROSE 3.3% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 30% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 40% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 5% AND ELECTROLYTE NO. 48 IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 DEXTROSE 5% AND POTASSIUM CHLORIDE 0.075%, DEXTROSE
 DEXTROSE 5% AND POTASSIUM CHLORIDE 0.075% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 5% AND POTASSIUM CHLORIDE 0.15%, DEXTROSE
 DEXTROSE 5% AND POTASSIUM CHLORIDE 0.15% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 5% AND POTASSIUM CHLORIDE 0.224% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 5% AND POTASSIUM CHLORIDE 0.3% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 5% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 5% AND SODIUM CHLORIDE 0.225%, DEXTROSE
 DEXTROSE 5% AND SODIUM CHLORIDE 0.225% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 5% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 5% AND SODIUM CHLORIDE 0.45%, DEXTROSE
 DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 5% AND SODIUM CHLORIDE 0.9%, DEXTROSE
 DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 5% IN LACTATED RINGER'S IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 5% IN SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 5% IN SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 5% IN SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 5% IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 10MEQ, DEXTROSE
 DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 15MEQ (K), DEXTROSE
 DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 20MEQ, DEXTROSE
 DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 20MEQ (K), DEXTROSE
 DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 30MEQ, DEXTROSE
 DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 40MEQ, DEXTROSE
 DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 5MEQ, DEXTROSE
 DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 5MEQ (K), DEXTROSE
 DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 10MEQ IN PLASTIC CONTAINER,
 DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 15MEQ IN PLASTIC CONTAINER,
 DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 20MEQ IN PLASTIC CONTAINER,
 DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 30MEQ IN PLASTIC CONTAINER,
 DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 40MEQ IN PLASTIC CONTAINER,
 DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 5MEQ IN PLASTIC CONTAINER,
 DEXTROSE 5%, SODIUM CHLORIDE 0.45% AND POTASSIUM CHLORIDE 20MEQ (K) IN PLASTIC CONTAINER,
 DEXTROSE 50%, DEXTROSE

APPENDIX A - PRODUCT NAME INDEX

** D **

DEXTROSE 50% IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 70% IN PLASTIC CONTAINER, DEXTROSE
DEXYCU KIT, DEXAMETHASONE
DHIVY, CARBIDOPA
DIABETA, GLYBURIDE
DIACOMIT, STIRIPENTOL
DIANEAL LOW CALCIUM W/ DEXTROSE 1.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
DIANEAL LOW CALCIUM W/ DEXTROSE 2.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
DIANEAL LOW CALCIUM W/ DEXTROSE 4.25% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
DIANEAL LOW CALCIUM W/ DEXTROSE 1.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
DIANEAL LOW CALCIUM W/ DEXTROSE 2.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
DIANEAL LOW CALCIUM W/ DEXTROSE 3.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
DIANEAL LOW CALCIUM W/ DEXTROSE 4.25% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
DIANEAL PD-2 W/ DEXTROSE 1.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
DIANEAL PD-2 W/ DEXTROSE 2.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
DIANEAL PD-2 W/ DEXTROSE 4.25% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
DIASTAT, DIAZEPAM
DIASTAT ACUDIAL, DIAZEPAM
DIATRIZOATE MEGLUMINE AND DIATRIZOATE SODIUM, DIATRIZOATE MEGLUMINE
DIAZEPAM, DIAZEPAM
DIAZEPAM INTENSOL, DIAZEPAM
DIAZOXIDE, DIAZOXIDE
DIBENZYLIN, PHENOXYBENZAMINE HYDROCHLORIDE
DICLEGIS, DOXYLAMINE SUCCINATE
DICLOFENAC POTASSIUM, DICLOFENAC POTASSIUM
DICLOFENAC SODIUM, DICLOFENAC SODIUM (OTC)
DICLOFENAC SODIUM, DICLOFENAC SODIUM
DICLOFENAC SODIUM AND MISOPROSTOL, DICLOFENAC SODIUM
DICLOXACILLIN SODIUM, DICLOXACILLIN SODIUM
DICYCLOMINE HYDROCHLORIDE, DICYCLOMINE HYDROCHLORIDE
DICYCLOMINE HYDROCHLORIDE (PRESERVATIVE FREE), DICYCLOMINE HYDROCHLORIDE
DIETHYLPROPION HYDROCHLORIDE, DIETHYLPROPION HYDROCHLORIDE
DIFFERIN, ADAPALENE (OTC)
DIFFERIN, ADAPALENE
DIFICID, FIDAXOMICIN
DIFLORASONE DIACETATE, DIFLORASONE DIACETATE
DIFLUCAN, FLUCONAZOLE
DIFLUNISAL, DIFLUNISAL
DIFLUPREDNATE, DIFLUPREDNATE
DIGOXIN, DIGOXIN
DIHYDROERGOTAMINE MESYLATE, DIHYDROERGOTAMINE MESYLATE
DILANTIN, PHENYTOIN
DILANTIN, PHENYTOIN SODIUM
DILANTIN-125, PHENYTOIN
DILAUDID, HYDROMORPHONE HYDROCHLORIDE
DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
DIMENHYDRINATE, DIMENHYDRINATE
DIMETHYL FUMARATE, DIMETHYL FUMARATE
DIOVAN, VALSARTAN
DIOVAN HCT, HYDROCHLOROTHIAZIDE
DIPENTUM, OLSALAZINE SODIUM
DIPHENHYDRAMINE HYDROCHLORIDE, DIPHENHYDRAMINE HYDROCHLORIDE
DIPHENHYDRAMINE HYDROCHLORIDE PRESERVATIVE FREE, DIPHENHYDRAMINE HYDROCHLORIDE
DIPHENOXYLATE HYDROCHLORIDE AND ATROPINE SULFATE, ATROPINE SULFATE
DIPRIVAN, PROPOFOL
DIPROLENE, BETAMETHASONE DIPROPIONATE
DIPYRIDAMOLE, DIPYRIDAMOLE
DISOPYRAMIDE PHOSPHATE, DISOPYRAMIDE PHOSPHATE
DISULFIRAM, DISULFIRAM
DIURIL, CHLOROTHIAZIDE
DIVALPROEX SODIUM, DIVALPROEX SODIUM
DIVIGEL, ESTRADIOL
DOBUTAMINE HYDROCHLORIDE, DOBUTAMINE HYDROCHLORIDE
DOBUTAMINE HYDROCHLORIDE IN DEXTROSE 5% IN PLASTIC CONTAINER, DOBUTAMINE HYDROCHLORIDE
DOCETAXEL, DOCETAXEL

APPENDIX A - PRODUCT NAME INDEX

** D **

DOCIVYX, DOCETAXEL
DOCOSANOL, DOCOSANOL (OTC)
DODEX, CYANOCOBALAMIN
DOFETILDE, DOFETILIDE
DOFETILIDE, DOFETILIDE
DOJOLVI, TRIHEPTANOIN
DOLISHALE, ETHINYL ESTRADIOL
DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
DOPAMINE HYDROCHLORIDE, DOPAMINE HYDROCHLORIDE
DOPAMINE HYDROCHLORIDE IN DEXTROSE 5% IN PLASTIC CONTAINER, DOPAMINE HYDROCHLORIDE
DOPRAM, DOXAPRAM HYDROCHLORIDE
DOPTelet, AVATROMBOPAG MALEATE
DORAL, QUAZEPAM
DORYX, DOXYCYCLINE HYCLATE
DORYX MPC, DOXYCYCLINE HYCLATE
DORZOLAMIDE HYDROCHLORIDE, DORZOLAMIDE HYDROCHLORIDE
DORZOLAMIDE HYDROCHLORIDE AND TIMOLOL MALEATE, DORZOLAMIDE HYDROCHLORIDE
DOTAREM, GADOTERATE MEGLUMINE
DOVATO, DOLUTEGRAVIR SODIUM
DOVONEX, CALCIPOTRIENE
DOXAPRAM HYDROCHLORIDE, DOXAPRAM HYDROCHLORIDE
DOXAZOSIN MESYLATE, DOXAZOSIN MESYLATE
DOXEPIN HYDROCHLORIDE, DOXEPIN HYDROCHLORIDE
DOXERCALCIFEROL, DOXERCALCIFEROL
DOXIL (LIPOSOMAL), DOXORUBICIN HYDROCHLORIDE
DOXORUBICIN HYDROCHLORIDE, DOXORUBICIN HYDROCHLORIDE
DOXORUBICIN HYDROCHLORIDE (LIPOSOMAL), DOXORUBICIN HYDROCHLORIDE
DOXY 100, DOXYCYCLINE HYCLATE
DOXY 200, DOXYCYCLINE HYCLATE
DOXYCYCLINE, DOXYCYCLINE
DOXYCYCLINE, DOXYCYCLINE HYCLATE
DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
DOXYLAMINE SUCCINATE, DOXYLAMINE SUCCINATE (OTC)
DOXYLAMINE SUCCINATE AND PYRIDOXINE HYDROCHLORIDE, DOXYLAMINE SUCCINATE
DRAX EXAMETAZIME, TECHNETIUM TC-99M EXAMETAZIME KIT
DRAXIMAGE DTPA, TECHNETIUM TC-99M PENTETATE KIT
DRAXIMAGE MDP-25, TECHNETIUM TC-99M MEDRONATE
DRISDOL, ERGOCALCIFEROL
DRIZALMA SPRINKLE, DULOXETINE HYDROCHLORIDE
DRONABINOL, DRONABINOL
DRONEDARONE HYDROCHLORIDE, DRONEDARONE HYDROCHLORIDE
DROPERIDOL, DROPERIDOL
DROSPIRENONE AND ETHINYL ESTRADIOL, DROSPIRENONE
DROSPIRENONE, ETHINYL ESTRADIOL AND LEVOMEFOLATE CALCIUM, DROSPIRENONE
DROXIDOPA, DROXIDOPA
DUAC, BENZOYL PEROXIDE
DUAKLIR PRESSAIR, ACLIDINIUM BROMIDE
DUAVEE, BAZEDOXIFENE ACETATE
DUETACT, GLIMEPIRIDE
DULERA, FORMOTEROL FUMARATE
DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE
DUOBRII, HALOBETASOL PROPIONATE
DUODOTE, ATROPINE
DUOPA, CARBIDOPA
DURACLON, CLONIDINE HYDROCHLORIDE
DURAMORPH PF, MORPHINE SULFATE
DURAPREP, IODINE POVACRYLEX (OTC)
DUREZOL, DIFLUPREDNATE
DURYSTA, BIMATOPROST
DUTASTERIDE, DUTASTERIDE
DUTASTERIDE AND TAMSULOSIN HYDROCHLORIDE, DUTASTERIDE
DUVOID, BETHANECHOL CHLORIDE
DUVYZAT, GIVINOSTAT HYDROCHLORIDE
DYANAVEL XR, AMPHETAMINE
DYANAVEL XR 10, AMPHETAMINE

APPENDIX A - PRODUCT NAME INDEX

** D **

DYANAVEL XR 15, AMPHETAMINE
 DYANAVEL XR 20, AMPHETAMINE
 DYANAVEL XR 5, AMPHETAMINE
 DYCLOPRO, DYCLONINE HYDROCHLORIDE
 DYMISTA, AZELASTINE HYDROCHLORIDE
 DYRENIUM, TRIAMTERENE

** E **

E-Z-HD, BARIUM SULFATE
 E-Z-PAQUE, BARIUM SULFATE
 E.E.S., ERYTHROMYCIN ETHYLSUCCINATE
 E.E.S. 400, ERYTHROMYCIN ETHYLSUCCINATE
 EC-NAPROSYN, NAPROXEN
 ECONAZOLE NITRATE, ECONAZOLE NITRATE
 ECOZA, ECONAZOLE NITRATE
 EDARAVONE, EDARAVONE
 EDARBI, AZILSARTAN KAMEDOXOMIL
 EDARBYCLOR, AZILSARTAN KAMEDOXOMIL
 EDECRIN, ETHACRYNATE SODIUM
 EDECRIN, ETHACRYNIC ACID
 EDETATE CALCIUM DISODIUM, EDETATE CALCIUM DISODIUM
 EDEX, ALPROSTADIL
 EDLUAR, ZOLPIDEM TARTRATE
 EDURANT, RILPIVIRINE HYDROCHLORIDE
 EFAVIRENZ, EFAVIRENZ
 EFAVIRENZ, EMTRICITABINE, AND TENOFOVIR DISOPROXIL FUMARATE, EFAVIRENZ
 EFAVIRENZ, LAMIVUDINE AND TENOFOVIR DISOPROXIL FUMARATE, EFAVIRENZ
 EFFEXOR XR, VENLAFAXINE HYDROCHLORIDE
 EFFIENT, PRASUGREL HYDROCHLORIDE
 EFINACONAZOLE, EFINACONAZOLE
 EFUDEX, FLUOROURACIL
 EGATEN, TRICLABENDAZOLE
 ELCYS, CYSTEINE HYDROCHLORIDE
 ELESTRIN, ESTRADIOL
 ELETRIPTAN HYDROBROMIDE, ELETRIPTAN HYDROBROMIDE
 ELIDEL, PIMECROLIMUS
 ELIGARD KIT, LEUPROLIDE ACETATE
 ELIGLUSTAT TARTRATE, ELIGLUSTAT TARTRATE
 ELIMITE, PERMETHRIN
 ELINEST, ETHINYL ESTRADIOL
 ELIQUIS, APIXABAN
 ELIXOPHYLLIN, THEOPHYLLINE
 ELLA, ULIPRISTAL ACETATE
 ELLENCE, EPIRUBICIN HYDROCHLORIDE
 ELLIOTTS B SOLUTION, CALCIUM CHLORIDE
 ELMIRON, PENTOSAN POLYSULFATE SODIUM
 ELTROMBOPAG OLAMINE, ELTROMBOPAG OLAMINE
 ELUCIREM, GADOPICLENOL
 ELURYNG, ETHINYL ESTRADIOL
 ELYXYB, CELECOXIB
 EMEND, APREPITANT
 EMEND, FOSAPREPITANT DIMEGLUMINE
 EMERPHED, EPHEDRINE SULFATE
 EMFLAZA, DEFLAZACORT
 EMPAVELI, PEGCETACOPLAN
 EMROSI, MINOCYCLINE HYDROCHLORIDE
 EMSAM, SELEGILINE
 EMTRICITABINE, EMTRICITABINE
 EMTRICITABINE AND TENOFOVIR DISOPROXIL FUMARATE, EMTRICITABINE
 EMTRICITABINE AND TENOFOVIR ALAFENAMIDE FUMARATE, EMTRICITABINE
 EMTRICITABINE AND TENOFOVIR DISOPROXIL FUMARATE, EMTRICITABINE
 EMTRIVA, EMTRICITABINE
 EMVERM, MEBENDAZOLE
 EMZAHH, NORETHINDRONE
 ENALAPRIL MALEATE, ENALAPRIL MALEATE

APPENDIX A - PRODUCT NAME INDEX

** E **

ENALAPRIL MALEATE AND HYDROCHLOROTHIAZIDE, ENALAPRIL MALEATE
ENALAPRILAT, ENALAPRILAT
ENDARI, L-GLUTAMINE
ENDOMETRIN, PROGESTERONE
ENILLORING, ETHINYL ESTRADIOL
ENOXAPARIN SODIUM, ENOXAPARIN SODIUM
ENOXAPARIN SODIUM (PRESERVATIVE FREE), ENOXAPARIN SODIUM
ENPRESSE-28, ETHINYL ESTRADIOL
ENSACOVE, ENSARTINIB HYDROCHLORIDE
ENSKYCE, DESOGESTREL
ENSTILAR, BETAMETHASONE DIPROPIONATE
ENTACAPONE, ENTACAPONE
ENTECAVIR, ENTECAVIR
ENTERO VU 24%, BARIUM SULFATE
ENTOCORT EC, BUDESONIDE
ENTRESTO, SACUBITRIL
ENTRESTO SPRINKLE, SACUBITRIL
ENVARUSUS XR, TACROLIMUS
ENZALUTAMIDE, ENZALUTAMIDE
EOHILIA, BUDESONIDE
EOVIST, GADOXETATE DISODIUM
EPANED, ENALAPRIL MALEATE
EPCLUSA, SOFOSBUVIR
EPHEDRINE SULFATE, EPHEDRINE SULFATE
EPIDIOLEX, CANNABIDIOL
EPIDUO, ADAPALENE
EPIDUO FORTE, ADAPALENE
EPIFOAM, HYDROCORTISONE ACETATE
EPINASTINE HYDROCHLORIDE, EPINASTINE HYDROCHLORIDE
EPINEPHRINE, EPINEPHRINE
EPINEPHRINE (AUTOINJECTOR), EPINEPHRINE
EPIPEN, EPINEPHRINE
EPIPEN JR., EPINEPHRINE
EPIRUBICIN HYDROCHLORIDE, EPIRUBICIN HYDROCHLORIDE
EPITOL, CARBAMAZEPINE
EPIVIR, LAMIVUDINE
EPLERENONE, EPLERENONE
EPOPROSTENOL SODIUM, EPOPROSTENOL SODIUM
EPRONTIA, TOPIRAMATE
EPSOLAY, BENZOYL PEROXIDE
EPTIFIBATIDE, EPTIFIBATIDE
EQUETRO, CARBAMAZEPINE
ERAXIS, ANIDULAFUNGIN
ERGOCALCIFEROL, ERGOCALCIFEROL
ERGOMAR, ERGOTAMINE TARTRATE
ERGOTAMINE TARTRATE AND CAFFEINE, CAFFEINE
ERIBULIN MESYLATE, ERIBULIN MESYLATE
ERIVEDGE, VISMODEGIB
ERLEADA, APALUTAMIDE
ERLOTINIB HYDROCHLORIDE, ERLOTINIB HYDROCHLORIDE
ERMEZA, LEVOTHYROXINE SODIUM
ERRIN, NORETHINDRONE
ERTACZO, SERTACONAZOLE NITRATE
ERTAPENEM SODIUM, ERTAPENEM SODIUM
ERTUGLIFLOZIN, ERTUGLIFLOZIN
ERY-TAB, ERYTHROMYCIN
ERYC, ERYTHROMYCIN
ERYGEL, ERYTHROMYCIN
ERYPED, ERYTHROMYCIN ETHYLSUCCINATE
ERYTHRA-DERM, ERYTHROMYCIN
ERYTHROCIN, ERYTHROMYCIN LACTOBIONATE
ERYTHROCIN STEARATE, ERYTHROMYCIN STEARATE
ERYTHROMYCIN, ERYTHROMYCIN
ERYTHROMYCIN AND BENZOYL PEROXIDE, BENZOYL PEROXIDE
ERYTHROMYCIN ETHYLSUCCINATE, ERYTHROMYCIN ETHYLSUCCINATE

APPENDIX A - PRODUCT NAME INDEX

** E **

ERYTHROMYCIN LACTOBIONATE, ERYTHROMYCIN LACTOBIONATE
 ERZOFRI, PALIPERIDONE PALMITATE
 ESBRIET, PIRFENIDONE
 ESCITALOPRAM OXALATE, ESCITALOPRAM OXALATE
 ESLICARBAZEPINE ACETATE, ESLICARBAZEPINE ACETATE
 ESMOLOL HYDROCHLORIDE, ESMOLOL HYDROCHLORIDE
 ESMOLOL HYDROCHLORIDE DOUBLE STRENGTH IN PLASTIC CONTAINER, ESMOLOL HYDROCHLORIDE
 ESMOLOL HYDROCHLORIDE IN PLASTIC CONTAINER, ESMOLOL HYDROCHLORIDE
 ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM (OTC)
 ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM
 ESOMEPRAZOLE SODIUM, ESOMEPRAZOLE SODIUM
 ESTARYLLA, ETHINYL ESTRADIOL
 ESTAZOLAM, ESTAZOLAM
 ESTRACE, ESTRADIOL
 ESTRADIOL, ESTRADIOL
 ESTRADIOL AND NORETHINDRONE ACETATE, ESTRADIOL
 ESTRADIOL VALERATE, ESTRADIOL VALERATE
 ESTRING, ESTRADIOL
 ESTROGEL, ESTRADIOL
 ESZOPICLONE, ESZOPICLONE
 ETHACRYNATE SODIUM, ETHACRYNATE SODIUM
 ETHACRYNIC ACID, ETHACRYNIC ACID
 ETHAMBUTOL HYDROCHLORIDE, ETHAMBUTOL HYDROCHLORIDE
 ETHAMOLIN, ETHANOLAMINE OLEATE
 ETHINYL ESTRADIOL AND NORELGESTROMIN, ETHINYL ESTRADIOL
 ETHINYL ESTRADIOL; ETONOGESTREL, ETHINYL ESTRADIOL
 ETHOSUXIMIDE, ETHOSUXIMIDE
 ETHYNODIOL DIACETATE AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
 ETODOLAC, ETODOLAC
 ETOMIDATE, ETOMIDATE
 ETOPOPHOS PRESERVATIVE FREE, ETOPOSIDE PHOSPHATE
 ETOPOSIDE, ETOPOSIDE
 ETRAVIRINE, ETRAVIRINE
 EUCRISA, CRISABOROLE
 EURAX, CROTAMITON
 EUTHYROX, LEVOTHYROXINE SODIUM **
 EVAMIST, ESTRADIOL
 EVEKEO, AMPHETAMINE SULFATE
 EVEROLIMUS, EVEROLIMUS
 EVISTA, RALOXIFENE HYDROCHLORIDE
 EVOMELA, MELPHALAN HYDROCHLORIDE
 EVOTAZ, ATAZANAVIR SULFATE
 EVOXAC, CEVIMELINE HYDROCHLORIDE
 EVRYSDI, RISDIPLAM
 EXCEDRIN (MIGRAINE RELIEF), ACETAMINOPHEN (OTC)
 EXELDERM, SULCONAZOLE NITRATE
 EXELON, RIVASTIGMINE
 EXEM FOAM KIT, AIR POLYMER-TYPE A
 EXEMESTANE, EXEMESTANE
 EXENATIDE SYNTHETIC, EXENATIDE SYNTHETIC
 EXFORGE, AMLODIPINE BESYLATE
 EXFORGE HCT, AMLODIPINE BESYLATE
 EXIDINE, CHLORHEXIDINE GLUCONATE (OTC)
 EXJADE, DEFERASIROX
 EXONDYS 51, ETEPLIRSEN
 EXPAREL, BUPIVACAINE
 EXTENDED PHENYTOIN SODIUM, PHENYTOIN SODIUM
 EXTRANEAL, ICODEXTRIN
 EYSUVIS, LOTEPIREDNOL ETABONATE
 EZETIMIBE, EZETIMIBE
 EZETIMIBE AND SIMVASTATIN, EZETIMIBE

** F **

FABHALTA, IPTACOPAN HYDROCHLORIDE
 FABIOR, TAZAROTENE

APPENDIX A - PRODUCT NAME INDEX

** F **

FALLBACK SOLO, LEVONORGESTREL (OTC)
FALMINA, ETHINYL ESTRADIOL
FAMCICLOVIR, FAMCICLOVIR
FAMOTIDINE, FAMOTIDINE (OTC)
FAMOTIDINE, FAMOTIDINE
FAMOTIDINE PRESERVATIVE FREE, FAMOTIDINE
FAMOTIDINE PRESERVATIVE FREE IN PLASTIC CONTAINER, FAMOTIDINE
FAMOTIDINE, CALCIUM CARBONATE, AND MAGNESIUM HYDROXIDE, CALCIUM CARBONATE (OTC)
FAMOTIDINE; IBUPROFEN, FAMOTIDINE
FANAPT, ILOPERIDONE
FARXIGA, DAPAGLIFLOZIN
FASLODEX, FULVESTRANT
FEBUXOSTAT, FEBUXOSTAT
FELBAMATE, FELBAMATE
FELBATOL, FELBAMATE
FELDENE, PIROXICAM
FELODIPINE, FELODIPINE
FEMARA, LETROZOLE
FEMLYV, ETHINYL ESTRADIOL
FEMRING, ESTRADIOL ACETATE
FENOFIBRATE, FENOFIBRATE
FENOFIBRATE (MICRONIZED), FENOFIBRATE
FENOFIBRIC ACID, CHOLINE FENOFIBRATE
FENOPROFEN CALCIUM, FENOPROFEN CALCIUM
FENSOLVI KIT, LEUPROLIDE ACETATE
FENTANYL CITRATE, FENTANYL CITRATE
FENTANYL CITRATE PRESERVATIVE FREE, FENTANYL CITRATE
FENTANYL-100, FENTANYL
FENTANYL-12, FENTANYL
FENTANYL-25, FENTANYL
FENTANYL-37, FENTANYL
FENTANYL-50, FENTANYL
FENTANYL-62, FENTANYL
FENTANYL-75, FENTANYL
FENTANYL-87, FENTANYL
FERAHEME, FERUMOXYTOL
FERRIPROX, DEFERIPRONE
FERRLECIT, FERRIC OXYHYDROXIDE
FERUMOXYTOL, FERUMOXYTOL
FESOTERODINE FUMARATE, FESOTERODINE FUMARATE
FETROJA, CEFIDEROCOL SULFATE TOSYLATE
FETZIMA, LEVOMILNACIPRAN HYDROCHLORIDE
FEXINIDAZOLE, FEXINIDAZOLE
FEXOFENADINE HYDROCHLORIDE, FEXOFENADINE HYDROCHLORIDE (OTC)
FEXOFENADINE HYDROCHLORIDE, FEXOFENADINE HYDROCHLORIDE
FEXOFENADINE HYDROCHLORIDE ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
FEXOFENADINE HYDROCHLORIDE AND PSEUDOEPHEDRINE HYDROCHLORIDE, FEXOFENADINE HYDROCHLORIDE
FEXOFENADINE HYDROCHLORIDE HIVES, FEXOFENADINE HYDROCHLORIDE (OTC)
FILSPARI, SPARSENTAN
FILSUVEZ, BIRCH TRITERPENES
FINACEA, AZELAIC ACID
FINASTERIDE, FINASTERIDE
FINGOLIMOD HYDROCHLORIDE, FINGOLIMOD HYDROCHLORIDE
FINTEPLA, FENFLURAMINE HYDROCHLORIDE
FINZALA, ETHINYL ESTRADIOL
FIORICET W/ CODEINE, ACETAMINOPHEN
FIRAZYR, ICATIBANT ACETATE
FIRDAPSE, AMIFAMPRIDINE PHOSPHATE
FIRMAGON, DEGARELIX ACETATE
FIRVANQ KIT, VANCOMYCIN HYDROCHLORIDE
FLAC, FLUOCINOLONE ACETONIDE
FLAGYL, METRONIDAZOLE
FLAGYL I.V. RTU IN PLASTIC CONTAINER, METRONIDAZOLE
FLAREX, FLUOROMETHOLONE ACETATE
FLAVOXATE HYDROCHLORIDE, FLAVOXATE HYDROCHLORIDE

APPENDIX A - PRODUCT NAME INDEX

** F **

FLECAINIDE ACETATE, FLECAINIDE ACETATE
FLECTOR, DICLOFENAC EPOLAMINE
FLEQSUVY, BACLOFEN
FLOLAN, EPOPROSTENOL SODIUM
FLOLIPID, SIMVASTATIN
FLOMAX, TAMSULOSIN HYDROCHLORIDE
FLONASE ALLERGY RELIEF, FLUTICASONE PROPIONATE (OTC)
FLONASE SENSIMIST ALLERGY RELIEF, FLUTICASONE FUROATE (OTC)
FLOVENT DISKUS 100, FLUTICASONE PROPIONATE
FLOVENT DISKUS 250, FLUTICASONE PROPIONATE
FLOVENT DISKUS 50, FLUTICASONE PROPIONATE
FLOVENT HFA, FLUTICASONE PROPIONATE
FLOXURIDINE, FLOXURIDINE
FLUCINOLONE ACETONIDE, FLUOCINOLONE ACETONIDE
FLUCONAZOLE, FLUCONAZOLE
FLUCONAZOLE IN SODIUM CHLORIDE 0.9%, FLUCONAZOLE
FLUCONAZOLE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, FLUCONAZOLE
FLUCYTOSINE, FLUCYTOSINE
FLUDARABINE PHOSPHATE, FLUDARABINE PHOSPHATE
FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18
FLUDROCORTISONE ACETATE, FLUDROCORTISONE ACETATE
FLUMAZENIL, FLUMAZENIL
FLUNISOLIDE, FLUNISOLIDE
FLUOCINOLONE ACETONIDE, FLUOCINOLONE ACETONIDE
FLUOCINONIDE, FLUOCINONIDE
FLUOCINONIDE ACETONIDE, FLUOCINOLONE ACETONIDE
FLUOCINONIDE EMULSIFIED BASE, FLUOCINONIDE
FLUORESCEIN SODIUM, FLUORESCEIN SODIUM
FLUORESCEIN SODIUM AND BENOXINATE HYDROCHLORIDE, BENOXINATE HYDROCHLORIDE
FLUORESCITE, FLUORESCEIN SODIUM
FLUORODOPA F18, FLUORODOPA F-18
FLUOROMETHOLONE, FLUOROMETHOLONE
FLUOROURACIL, FLUOROURACIL
FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
FLUPHENAZINE DECANOATE, FLUPHENAZINE DECANOATE
FLUPHENAZINE HYDROCHLORIDE, FLUPHENAZINE HYDROCHLORIDE
FLURANDRENOLIDE, FLURANDRENOLIDE
FLURAZEPAM HYDROCHLORIDE, FLURAZEPAM HYDROCHLORIDE
FLURBIPROFEN, FLURBIPROFEN
FLURBIPROFEN SODIUM, FLURBIPROFEN SODIUM
FLUTAMIDE, FLUTAMIDE
FLUTICASONE PROPIONATE, FLUTICASONE PROPIONATE (OTC)
FLUTICASONE PROPIONATE, FLUTICASONE PROPIONATE
FLUTICASONE PROPIONATE AND SALMETEROL XINAFOATE, FLUTICASONE PROPIONATE
FLUVASTATIN SODIUM, FLUVASTATIN SODIUM
FLUVOXAMINE MALEATE, FLUVOXAMINE MALEATE
FLYRCADO, FLURPIRIDAZ F-18
FML, FLUOROMETHOLONE
FML FORTE, FLUOROMETHOLONE
FOCALIN, DEXMETHYLPHENIDATE HYDROCHLORIDE
FOCALIN XR, DEXMETHYLPHENIDATE HYDROCHLORIDE
FOCINVEZ, FOSAPREPITANT DIMEGLUMINE
FOLIC ACID, FOLIC ACID
FOLOTYN, PRALATREXATE
FOMEPIZOLE, FOMEPIZOLE
FONDAPARINUX SODIUM, FONDAPARINUX SODIUM
FORANE, ISOFLURANE
FORFIVO XL, BUPROPION HYDROCHLORIDE
FORMOTEROL FUMARATE, FORMOTEROL FUMARATE
FORTAMET, METFORMIN HYDROCHLORIDE
FORTEO, TERIPARATIDE
FOSAMAX, ALENDRONATE SODIUM
FOSAMAX PLUS D, ALENDRONATE SODIUM
FOSAMPRENAVIR CALCIUM, FOSAMPRENAVIR CALCIUM
FOSAPREPITANT DIMEGLUMINE, FOSAPREPITANT DIMEGLUMINE

APPENDIX A - PRODUCT NAME INDEX

** F **

FOSCARNET SODIUM, FOSCARNET SODIUM
 FOSCAVIR, FOSCARNET SODIUM
 FOSFOMYCIN TROMETHAMINE, FOSFOMYCIN TROMETHAMINE
 FOSINOPRIL SODIUM, FOSINOPRIL SODIUM
 FOSINOPRIL SODIUM AND HYDROCHLOROTHIAZIDE, FOSINOPRIL SODIUM
 FOSPHENYTOIN SODIUM, FOSPHENYTOIN SODIUM
 FOSRENOL, LANTHANUM CARBONATE
 FOTIVDA, TIVOZANIB HYDROCHLORIDE
 FRAGMIN, DALTEPARIN SODIUM
 FROVA, FROVATRIPTAN SUCCINATE
 FROVATRIPTAN SUCCINATE, FROVATRIPTAN SUCCINATE
 FRUZAQLA, FRUQUINTINIB
 FULVESTRANT, FULVESTRANT
 FULVICIN P/G, GRISEOFULVIN, ULTRAMICROSIZE
 FULVICIN P/G 165, GRISEOFULVIN, ULTRAMICROSIZE
 FULVICIN P/G 330, GRISEOFULVIN, ULTRAMICROSIZE
 FULVICIN-U/F, GRISEOFULVIN, MICROSIZE
 FURADANTIN, NITROFURANTOIN
 FUROSCIX, FUROSEMIDE
 FUROSEMIDE, FUROSEMIDE
 FUZEON, ENFUVRTIDE
 FYARRO, SIROLIMUS
 FYAVOLV, ETHINYL ESTRADIOL
 FYCOMPA, PERAMPANEL
 FYREMADEL, GANIRELIX ACETATE

** G **

GABAPENTIN, GABAPENTIN
 GABITRIL, TIAGABINE HYDROCHLORIDE
 GABLOFEN, BACLOFEN
 GADAVIST, GADOBUTROL
 GADOBUTROL, GADOBUTROL
 GADOTERATE MEGLUMINE, GADOTERATE MEGLUMINE
 GALAFOLD, MIGALASTAT HYDROCHLORIDE
 GALANTAMINE HYDROBROMIDE, GALANTAMINE HYDROBROMIDE
 GALLIUM CITRATE GA 67, GALLIUM CITRATE GA-67
 GALLIUM GA 68 EDOTREOTIDE, GALLIUM GA-68 EDOTREOTIDE
 GALLIUM GA 68 GOZETOTIDE, GALLIUM GA-68 GOZETOTIDE
 GALZIN, ZINC ACETATE
 GANCICLOVIR SODIUM, GANCICLOVIR SODIUM
 GANIRELIX ACETATE, GANIRELIX ACETATE
 GANZYK-RTU, GANCICLOVIR
 GASTROCROM, CROMOLYN SODIUM
 GASTROGRAFIN, DIATRIZOATE MEGLUMINE
 GATIFLOXACIN, GATIFLOXACIN
 GATTEX KIT, TEDUGLUTIDE
 GAVRETO, PRALSETINIB
 GEFITINIB, GEFITINIB
 GEMCITABINE HYDROCHLORIDE, GEMCITABINE HYDROCHLORIDE
 GEMFIBROZIL, GEMFIBROZIL
 GEMMILY, ETHINYL ESTRADIOL
 GEMTESA, VIBEGRON
 GENERLAC, LACTULOSE
 GENGRAF, CYCLOSPORINE
 GENOSYL, NITRIC OXIDE
 GENTAMICIN SULFATE, GENTAMICIN SULFATE
 GENTAMICIN SULFATE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, GENTAMICIN SULFATE
 GENVOYA, COBICISTAT
 GEODON, ZIPRASIDONE HYDROCHLORIDE
 GEODON, ZIPRASIDONE MESYLATE
 GIAPREZA, ANGIOTENSIN II ACETATE
 GILDAGIA, ETHINYL ESTRADIOL
 GILDESS 24 FE, ETHINYL ESTRADIOL
 GILENYA, FINGOLIMOD HYDROCHLORIDE
 GILOTRIF, AFATINIB DIMALEATE

APPENDIX A - PRODUCT NAME INDEX

** G **

GIMOTI, METOCLOPRAMIDE HYDROCHLORIDE
 GIVLAARI, GIVOSIRAN SODIUM
 GLATIRAMER ACETATE, GLATIRAMER ACETATE
 GLATOPA, GLATIRAMER ACETATE
 GLEEVEC, IMATINIB MESYLATE
 GLEOLAN, AMINOLEVULINIC ACID HYDROCHLORIDE
 GLEOSTINE, LOMUSTINE
 GLIADEL, CARMUSTINE
 GLIMEPIRIDE, GLIMEPIRIDE
 GLIPIZIDE, GLIPIZIDE
 GLIPIZIDE AND METFORMIN HYDROCHLORIDE, GLIPIZIDE
 GLOFIL-125, IOTHALAMATE SODIUM I-125
 GLOPERBA, COLCHICINE
 GLUCAGON, GLUCAGON
 GLUCAGON, GLUCAGON HYDROCHLORIDE
 GLUCOTROL XL, GLIPIZIDE
 GLUMETZA, METFORMIN HYDROCHLORIDE
 GLYBURIDE, GLYBURIDE
 GLYBURIDE (MICRONIZED), GLYBURIDE
 GLYBURIDE AND METFORMIN HYDROCHLORIDE, GLYBURIDE
 GLYCINE 1.5% IN PLASTIC CONTAINER, GLYCINE
 GLYCOPYRROLATE, GLYCOPYRROLATE
 GLYDO, LIDOCAINE HYDROCHLORIDE
 GLYNASE, GLYBURIDE
 GLYRX-PF, GLYCOPYRROLATE
 GLYSET, MIGLITOL
 GLYXAMBI, EMPAGLIFLOZIN
 GOCOVRI, AMANTADINE HYDROCHLORIDE
 GOLYTELY, POLYETHYLENE GLYCOL 3350
 GOPRELTO, COCAINE HYDROCHLORIDE
 GRALISE, GABAPENTIN
 GRANISETRON HYDROCHLORIDE, GRANISETRON HYDROCHLORIDE
 GRANISETRON HYDROCHLORIDE PRESERVATIVE FREE, GRANISETRON HYDROCHLORIDE
 GRIS-PEG, GRISEOFULVIN, ULTRAMICROSIZE
 GRISEOFULVIN, GRISEOFULVIN, MICROSIZE
 GRISEOFULVIN, ULTRAMICROSIZE, GRISEOFULVIN, ULTRAMICROSIZE
 GRISEOFULVIN, ULTRAMICROSIZE, GRISEOFULVIN, ULTRAMICROSIZE
 GUAIFENESIN, GUAIFENESIN (OTC)
 GUAIFENESIN AND DEXTROMETHORPHAN HYDROBROMIDE, DEXTROMETHORPHAN HYDROBROMIDE (OTC)
 GUAIFENESIN AND PSEUDOEPHEDRINE HYDROCHLORIDE, GUAIFENESIN (OTC)
 GUANFACINE HYDROCHLORIDE, GUANFACINE HYDROCHLORIDE
 GVOKE HYOPEN, GLUCAGON
 GVOKE KIT, GLUCAGON
 GVOKE PFS, GLUCAGON
 GYNAZOLE-1, BUTOCONAZOLE NITRATE

** H **

HABITROL, NICOTINE (OTC)
 HAILEY 1.5/30, ETHINYL ESTRADIOL
 HAILEY 24 FE, ETHINYL ESTRADIOL
 HAILEY FE 1.5/30, ETHINYL ESTRADIOL
 HAILEY FE 1/20, ETHINYL ESTRADIOL
 HALAVEN, ERIBULIN MESYLATE
 HALCINONIDE, HALCINONIDE
 HALCION, TRIAZOLAM
 HALDOL, HALOPERIDOL DECANOATE
 HALOBETASOL PROPIONATE, HALOBETASOL PROPIONATE
 HALOETTE, ETHINYL ESTRADIOL
 HALOG, HALCINONIDE
 HALOPERIDOL, HALOPERIDOL
 HALOPERIDOL, HALOPERIDOL LACTATE
 HALOPERIDOL DECANOATE, HALOPERIDOL DECANOATE
 HARVONI, LEDIPASVIR
 HEATHER, NORETHINDRONE
 HECTOROL, DOXERCALCIFEROL

APPENDIX A - PRODUCT NAME INDEX

** H **

HEMABATE, CARBOPROST TROMETHAMINE
HEMADY, DEXAMETHASONE
HEMANGEOL, PROPRANOLOL HYDROCHLORIDE
HEPARIN SODIUM, HEPARIN SODIUM
HEPARIN SODIUM 1,000 UNITS AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, HEPARIN SODIUM
HEPARIN SODIUM 1,000 UNITS IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, HEPARIN SODIUM
HEPARIN SODIUM 10,000 UNITS IN DEXTROSE 5% IN PLASTIC CONTAINER, HEPARIN SODIUM
HEPARIN SODIUM 12,500 UNITS IN DEXTROSE 5% IN PLASTIC CONTAINER, HEPARIN SODIUM
HEPARIN SODIUM 12,500 UNITS IN SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, HEPARIN SODIUM
HEPARIN SODIUM 2,000 UNITS AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, HEPARIN SODIUM
HEPARIN SODIUM 2,000 UNITS IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, HEPARIN SODIUM
HEPARIN SODIUM 20,000 UNITS IN DEXTROSE 5% IN PLASTIC CONTAINER, HEPARIN SODIUM
HEPARIN SODIUM 25,000 UNITS IN DEXTROSE 5%, HEPARIN SODIUM
HEPARIN SODIUM 25,000 UNITS IN DEXTROSE 5% IN PLASTIC CONTAINER, HEPARIN SODIUM
HEPARIN SODIUM 25,000 UNITS IN SODIUM CHLORIDE 0.45%, HEPARIN SODIUM
HEPARIN SODIUM 25,000 UNITS IN SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, HEPARIN SODIUM
HEPARIN SODIUM IN PLASTIC CONTAINER, HEPARIN SODIUM
HEPARIN SODIUM PRESERVATIVE FREE, HEPARIN SODIUM
HEPZATO, MELPHALAN HYDROCHLORIDE
HER STYLE, LEVONORGESTREL (OTC)
HETLIOZ, TASIMELTEON
HETLIOZ LQ, TASIMELTEON
HIBICLENS, CHLORHEXIDINE GLUCONATE (OTC)
HIBISTAT, CHLORHEXIDINE GLUCONATE (OTC)
HICON, SODIUM IODIDE I-131
HIPREX, METHENAMINE HIPPURATE
HOMATROPINE METHYLBROMIDE AND HYDROCODONE BITARTRATE, HOMATROPINE METHYLBROMIDE
HORIZANT, GABAPENTIN ENACARBIL
HYCANTIN, TOPOTECAN HYDROCHLORIDE
HYCODAN, HOMATROPINE METHYLBROMIDE
HYDRA-ZIDE, HYDRALAZINE HYDROCHLORIDE
HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
HYDREA, HYDROXYUREA
HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
HYDROCHLOROTHIAZIDE AND OLMESARTAN MEDOXOMIL, HYDROCHLOROTHIAZIDE
HYDROCODONE BITARTRATE, HYDROCODONE BITARTRATE
HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
HYDROCODONE BITARTRATE AND ASPIRIN, ASPIRIN
HYDROCODONE BITARTRATE AND HOMATROPINE METHYLBROMIDE, HOMATROPINE METHYLBROMIDE
HYDROCODONE BITARTRATE AND IBUPROFEN, HYDROCODONE BITARTRATE
HYDROCODONE BITARTRATE, CHLORPHENIRAMINE MALEATE AND PSEUDOEPHEDRINE HYDROCHLORIDE,
HYDROCODONE POLISTIREX AND CHLORPHENIRAMINE POLISTIREX, CHLORPHENIRAMINE POLISTIREX
HYDROCORTISONE, HYDROCORTISONE
HYDROCORTISONE AND ACETIC ACID, ACETIC ACID, GLACIAL
HYDROCORTISONE BUTYRATE, HYDROCORTISONE BUTYRATE
HYDROCORTISONE IN ABSORBASE, HYDROCORTISONE
HYDROCORTISONE SODIUM SUCCINATE, HYDROCORTISONE SODIUM SUCCINATE
HYDROCORTISONE VALERATE, HYDROCORTISONE VALERATE
HYDROMORPHONE HYDROCHLORIDE, HYDROMORPHONE HYDROCHLORIDE
HYDROXOCOBALAMIN, HYDROXOCOBALAMIN
HYDROXYCHLOROQUINE SULFATE, HYDROXYCHLOROQUINE SULFATE
HYDROXYUREA, HYDROXYUREA
HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE
HYDROXYZINE PAMOATE, HYDROXYZINE PAMOATE
HYFTOR, SIROLIMUS
HYSINGLA ER, HYDROCODONE BITARTRATE
HYZAAR, HYDROCHLOROTHIAZIDE

** I **

IBANDRONATE SODIUM, IBANDRONATE SODIUM
IBRANCE, PALBOCICLIB
IBSRELA, TENAPANOR HYDROCHLORIDE
IBUPROFEN, IBUPROFEN (OTC)
IBUPROFEN, IBUPROFEN
IBUPROFEN AND DIPHENHYDRAMINE CITRATE, DIPHENHYDRAMINE CITRATE (OTC)

APPENDIX A - PRODUCT NAME INDEX

** I **

IBUPROFEN AND DIPHENHYDRAMINE HYDROCHLORIDE, DIPHENHYDRAMINE HYDROCHLORIDE (OTC)
IBUPROFEN AND FAMOTIDINE, FAMOTIDINE
IBUPROFEN AND PHENYLEPHRINE HYDROCHLORIDE, IBUPROFEN (OTC)
IBUPROFEN AND PSEUDOEPHEDRINE HYDROCHLORIDE, IBUPROFEN (OTC)
IBUPROFEN LYSINE, IBUPROFEN LYSINE
IBUPROHM COLD AND SINUS, IBUPROFEN (OTC)
IBUTILIDE FUMARATE, IBUTILIDE FUMARATE
ICATIBANT ACETATE, ICATIBANT ACETATE
ICLEVIA, ETHINYL ESTRADIOL
ICLUSIG, PONATINIB HYDROCHLORIDE
ICOSAPENT ETHYL, ICOSAPENT ETHYL
IDAMYCIN PFS, IDARUBICIN HYDROCHLORIDE
IDARUBICIN HYDROCHLORIDE, IDARUBICIN HYDROCHLORIDE
IDHIFA, ENASIDENIB MESYLATE
IDKIT:HP, CITRIC ACID
IDOSE TR, TRAVOPROST
IFEX, IFOSFAMIDE
IFOSFAMIDE, IFOSFAMIDE
IGALMI, DEXMEDETOMIDINE HYDROCHLORIDE
IHEEZO, CHLOROPROCAINE HYDROCHLORIDE
ILEVRO, NEPAFENAC
ILLUCCIX, GALLIUM GA-68 GOZETOTIDE
ILUVIEN, FLUOCINOLONE ACETONIDE
IMATINIB MESYLATE, IMATINIB MESYLATE
IMBRUVICA, IBRUTINIB
IMCIVREE, SETMELANOTIDE ACETATE
IMIPENEM AND CILASTATIN, CILASTATIN SODIUM
IMIPRAMINE HYDROCHLORIDE, IMIPRAMINE HYDROCHLORIDE
IMIPRAMINE PAMOATE, IMIPRAMINE PAMOATE
IMIQUIMOD, IMIQUIMOD
IMITREX, SUMATRIPTAN
IMITREX, SUMATRIPTAN SUCCINATE
IMITREX STATDOSE, SUMATRIPTAN SUCCINATE
IMKELDI, IMATINIB MESYLATE
IMMPHENTIV, PHENYLEPHRINE HYDROCHLORIDE
IMODIUM A-D, LOPERAMIDE HYDROCHLORIDE (OTC)
IMODIUM MULTI-SYMPTOM RELIEF, LOPERAMIDE HYDROCHLORIDE (OTC)
IMPAVIDO, MILTEFOSINE
IMPOYZ, CLOBETASOL PROPIONATE
IMURAN, AZATHIOPRINE
IMVEXXY, ESTRADIOL
INBRIJA, LEVODOPA
INCASSIA, NORETHINDRONE
INCRUSE ELLIPTA, UMECLIDINIUM BROMIDE
INDAPAMIDE, INDAPAMIDE
INDERAL LA, PROPRANOLOL HYDROCHLORIDE
INDIUM IN 111 OXYQUINOLINE, INDIUM IN-111 OXYQUINOLINE
INDIUM IN-111 PENTETREOTIDE KIT, INDIUM IN-111 PENTETREOTIDE KIT
INDOCIN, INDOMETHACIN
INDOCYANINE GREEN, INDOCYANINE GREEN
INDOMETHACIN, INDOMETHACIN
INFANT'S ADVIL, IBUPROFEN (OTC)
INFANTS' FEVERALL, ACETAMINOPHEN (OTC)
INFED, FERRIC OXYHYDROXIDE
INFUMORPH, MORPHINE SULFATE
INFUVITE ADULT, ALPHA-TOCOPHEROL ACETATE
INFUVITE PEDIATRIC, ASCORBIC ACID
INFUVITE PEDIATRIC (PHARMACY BULK PACKAGE), ASCORBIC ACID
INGREZZA, VALBENAZINE TOSYLATE
INGREZZA SPRINKLE, VALBENAZINE TOSYLATE
INJECTAFER, FERRIC CARBOXYMALTOSE
INLYTA, AXITINIB
INNOPRAN XL, PROPRANOLOL HYDROCHLORIDE
INOMAX, NITRIC OXIDE
INPEFA, SOTAGLIFLOZIN

APPENDIX A - PRODUCT NAME INDEX

** I **

INQOVI, CEDAZURIDINE
INREBIC, FEDRATINIB HYDROCHLORIDE
INSPRA, EPLERENONE
INTELENCE, ETRAVIRINE
INTRALIPID 20%, SOYBEAN OIL
INTRALIPID 30%, SOYBEAN OIL
INTRAROSA, PRASTERONE
INTROVALE, ETHINYL ESTRADIOL
INTUNIV, GUANFACINE HYDROCHLORIDE
INVANZ, ERTAPENEM SODIUM
INVEGA, PALIPERIDONE
INVEGA HAFYERA, PALIPERIDONE PALMITATE
INVEGA SUSTENNA, PALIPERIDONE PALMITATE
INVEGA TRINZA, PALIPERIDONE PALMITATE
INVELTYS, LOTEHPREDNOL ETABONATE
INVOKAMET, CANAGLIFLOZIN
INVOKAMET XR, CANAGLIFLOZIN
INVOKANA, CANAGLIFLOZIN
IOBENGUANE I-123, IOBENGUANE SULFATE I-123
IODIXANOL, IODIXANOL
IOFLUPANE I-123, IOFLUPANE I-123
IOMERVU, IOMEPROL
IONOSOL MB AND DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
IOPAMIDOL, IOPAMIDOL
IOPIDINE, APRACLONIDINE HYDROCHLORIDE
IOSAT, POTASSIUM IODIDE (OTC)
IPRATROPIUM BROMIDE, IPRATROPIUM BROMIDE
IQIRVO, ELAFIBRANOR
IRBESARTAN, IRBESARTAN
IRBESARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
IRESSA, GEFITINIB
IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE
ISENTRESS, RALTEGRAVIR POTASSIUM
ISENTRESS HD, RALTEGRAVIR POTASSIUM
ISIBLOOM, DESOGESTREL
ISOFLURANE, ISOFLURANE
ISOLYTE P IN DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
ISOLYTE S IN PLASTIC CONTAINER, MAGNESIUM CHLORIDE
ISOLYTE S PH 7.4 IN PLASTIC CONTAINER, MAGNESIUM CHLORIDE
ISONIAZID, ISONIAZID
ISOPROTERENOL HYDROCHLORIDE, ISOPROTERENOL HYDROCHLORIDE
ISOPTO ATROPINE, ATROPINE SULFATE
ISOPTO CARPINE, PILOCARPINE HYDROCHLORIDE
ISORDIL, ISOSORBIDE DINITRATE
ISOSORBIDE DINITRATE, ISOSORBIDE DINITRATE
ISOSORBIDE DINITRATE AND HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
ISOSORBIDE MONONITRATE, ISOSORBIDE MONONITRATE
ISOSULFAN BLUE, ISOSULFAN BLUE
ISOTRETINOIN, ISOTRETINOIN
ISOVUE-200, IOPAMIDOL
ISOVUE-250, IOPAMIDOL
ISOVUE-300, IOPAMIDOL
ISOVUE-370, IOPAMIDOL
ISOVUE-M 200, IOPAMIDOL
ISOVUE-M 300, IOPAMIDOL
ISRADIPINE, ISRADIPINE
ISTALOL, TIMOLOL MALEATE
ISTODAX, ROMIDEPSIN
ISTURISA, OSILODROSTAT PHOSPHATE
ITOVEBI, INAVOLISIB
ITRACONAZOLE, ITRACONAZOLE
IVABRADINE HYDROCHLORIDE, IVABRADINE HYDROCHLORIDE
IVERMECTIN, IVERMECTIN (OTC)
IVERMECTIN, IVERMECTIN
IVRA, MELPHALAN HYDROCHLORIDE

APPENDIX A - PRODUCT NAME INDEX

** I **

IVY BLOCK, BENTOQUATAM (OTC)
 IWILFIN, EFLORNITHINE HYDROCHLORIDE
 IXEMPRA KIT, IXABEPILONE
 IYUZEH, LATANOPROST
 IZERVAY, AVACINCAPTAD PEGOL SODIUM

** J **

JADENU, DEFERASIROX
 JADENU SPRINKLE, DEFERASIROX
 JAIMIESS, ETHINYL ESTRADIOL
 JAKAFI, RUXOLITINIB PHOSPHATE
 JALYN, DUTASTERIDE
 JANTOVEN, WARFARIN SODIUM
 JANUMET, METFORMIN HYDROCHLORIDE
 JANUMET XR, METFORMIN HYDROCHLORIDE
 JANUVIA, SITAGLIPTIN PHOSPHATE
 JARDIANCE, EMPAGLIFLOZIN
 JATENZO, TESTOSTERONE UNDECANOATE
 JAYPIRCA, PIRTOBRUTINIB
 JELMYTO, MITOMYCIN
 JENCYCLA, NORETHINDRONE
 JENTADUETO, LINAGLIPTIN
 JENTADUETO XR, LINAGLIPTIN
 JEVTANA KIT, CABAZITAXEL
 JOENJA, LENIOLISIB PHOSPHATE
 JORNAY PM, METHYLPHENIDATE HYDROCHLORIDE
 JUBLIA, EFINACONAZOLE
 JULUCA, DOLUTEGRAVIR SODIUM
 JUNEL 1.5/30, ETHINYL ESTRADIOL
 JUNEL 1/20, ETHINYL ESTRADIOL
 JUNEL FE 1.5/30, ETHINYL ESTRADIOL
 JUNEL FE 1/20, ETHINYL ESTRADIOL
 JUNIOR STRENGTH ADVIL, IBUPROFEN (OTC)
 JUNIOR STRENGTH MOTRIN, IBUPROFEN (OTC)
 JUXTAPID, LOMITAPIDE MESYLATE
 JYLAMVO, METHOTREXATE
 JYNARQUE, TOLVAPTAN

** K **

KABIVEN IN PLASTIC CONTAINER, AMINO ACIDS
 KAITLIB FE, ETHINYL ESTRADIOL
 KALETRA, LOPINAVIR
 KALEXATE, SODIUM POLYSTYRENE SULFONATE
 KALLIGA, DESOGESTREL
 KALYDECO, IVACAFTOR
 KAPSPARGO SPRINKLE, METOPROLOL SUCCINATE
 KARBINAL ER, CARBINOXAMINE MALEATE
 KARIVA, DESOGESTREL
 KATERZIA, AMLODIPINE BENZOATE
 KAZANO, ALOGLIPTIN BENZOATE
 KELNOR, ETHINYL ESTRADIOL
 KENALOG-10, TRIAMCINOLONE ACETONIDE
 KENALOG-40, TRIAMCINOLONE ACETONIDE
 KENALOG-80, TRIAMCINOLONE ACETONIDE
 KENGREAL, CANGRELOR
 KEPPRA, LEVETIRACETAM
 KEPPRA XR, LEVETIRACETAM
 KERENDIA, FINERENONE
 KETALAR, KETAMINE HYDROCHLORIDE
 KETAMINE HYDROCHLORIDE, KETAMINE HYDROCHLORIDE
 KETOCONAZOLE, KETOCONAZOLE
 KETOPROFEN, KETOPROFEN
 KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
 KETOROLAC TROMETHAMINE AND PHENYLEPHRINE HYDROCHLORIDE, KETOROLAC TROMETHAMINE
 KETOTIFEN FUMARATE, KETOTIFEN FUMARATE (OTC)

APPENDIX A - PRODUCT NAME INDEX

** K **

KETOZOLE, KETOCONAZOLE
 KEVEYIS, DICHLORPHENAMIDE
 KHAPZORY, LEVOLEUCOVORIN
 KIMYRSA, ORITAVANCIN DIPHOSPHATE
 KINEVAC, SINCALIDE
 KIONEX, SODIUM POLYSTYRENE SULFONATE
 KISQALI, RIBOCICLIB SUCCINATE
 KISQALI FEMARA CO-PACK (COPACKAGED), LETROZOLE
 KITABIS PAK, TOBRAMYCIN
 KLARON, SULFACETAMIDE SODIUM
 KLISYRI, TIRBANIBULIN
 KLONOPIN, CLONAZEPAM
 KLOR-CON, POTASSIUM CHLORIDE
 KLOR-CON M10, POTASSIUM CHLORIDE
 KLOR-CON M15, POTASSIUM CHLORIDE
 KLOR-CON M20, POTASSIUM CHLORIDE
 KLOXXADO, NALOXONE HYDROCHLORIDE
 KONVOMEK, OMEPRAZOLE
 KORLYM, MIFEPRISTONE
 KORSUVA, DIFELIKEFALIN ACETATE
 KOSELUGO, SELUMETINIB SULFATE
 KOVANAZE, OXYMETAZOLINE HYDROCHLORIDE
 KRAZATI, ADAGRASIB
 KRINTAFEL, TAFENOQUINE SUCCINATE
 KURVELO, ETHINYL ESTRADIOL
 KUVAN, SAPROPTERIN DIHYDROCHLORIDE
 KYBELLA, DEOXYCHOLIC ACID
 KYLEENA, LEVONORGESTREL
 KYPROLIS, CARFILZOMIB
 KYZATREX, TESTOSTERONE UNDECANOATE

** L **

L-GLUTAMINE, L-GLUTAMINE
 LABETALOL HYDROCHLORIDE, LABETALOL HYDROCHLORIDE
 LABETALOL HYDROCHLORIDE IN DEXTROSE, LABETALOL HYDROCHLORIDE
 LABETALOL HYDROCHLORIDE IN SODIUM CHLORIDE, LABETALOL HYDROCHLORIDE
 LACOSAMIDE, LACOSAMIDE
 LACTATED RINGER'S AND DEXTROSE 5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 LACTATED RINGER'S IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 LACTULOSE, LACTULOSE
 LAMICTAL, LAMOTRIGINE
 LAMICTAL CD, LAMOTRIGINE
 LAMICTAL ODT, LAMOTRIGINE
 LAMICTAL XR, LAMOTRIGINE
 LAMISIL, TERBINAFINE HYDROCHLORIDE (OTC)
 LAMISIL AT, TERBINAFINE (OTC)
 LAMISIL AT, TERBINAFINE HYDROCHLORIDE (OTC)
 LAMIVUDINE, LAMIVUDINE
 LAMIVUDINE AND ZIDOVUDINE, LAMIVUDINE
 LAMOTRIGINE, LAMOTRIGINE
 LAMPIT, NIFURTIMOX
 LANORINAL, ASPIRIN
 LANOXIN, DIGOXIN
 LANOXIN PEDIATRIC, DIGOXIN
 LANREOTIDE ACETATE, LANREOTIDE ACETATE
 LANSOPRAZOLE, LANSOPRAZOLE (OTC)
 LANSOPRAZOLE, LANSOPRAZOLE
 LANSOPRAZOLE, AMOXICILLIN AND CLARITHROMYCIN (COPACKAGED), AMOXICILLIN
 LANTHANUM CARBONATE, LANTHANUM CARBONATE
 LAPATINIB DITOSYLATE, LAPATINIB DITOSYLATE
 LARIN 1.5/30, ETHINYL ESTRADIOL
 LARIN 1/20, ETHINYL ESTRADIOL
 LARIN 24 FE, ETHINYL ESTRADIOL
 LARIN FE 1.5/30, ETHINYL ESTRADIOL
 LARIN FE 1/20, ETHINYL ESTRADIOL

APPENDIX A - PRODUCT NAME INDEX

** L **

LAROTID, AMOXICILLIN
 LARYNG-O-JET KIT, LIDOCAINE HYDROCHLORIDE
 LASIX, FUROSEMIDE
 LASTACFT, ALCAFTADINE (OTC)
 LATANOPROST, LATANOPROST
 LATISSE, BIMATOPROST
 LATUDA, LURASIDONE HYDROCHLORIDE
 LAZCLUZE, LAZERTINIB MESYLATE
 LEFLUNOMIDE, LEFLUNOMIDE
 LENALIDOMIDE, LENALIDOMIDE
 LENVIMA, LENVATINIB MESYLATE
 LEQSELVI, DEURUXOLITINIB PHOSPHATE
 LEQVIO, INCLISIRAN SODIUM
 LERIBANE, ETHINYL ESTRADIOL
 LESCOL XL, FLUVASTATIN SODIUM
 LESSINA-28, ETHINYL ESTRADIOL
 LETAIRIS, AMBRISENTAN
 LETROZOLE, LETROZOLE
 LEUCOVORIN CALCIUM, LEUCOVORIN CALCIUM
 LEUCOVORIN CALCIUM PRESERVATIVE FREE, LEUCOVORIN CALCIUM
 LEUKERAN, CHLORAMBUCIL
 LEUPROLIDE ACETATE, LEUPROLIDE ACETATE
 LEUPROLIDE ACETATE FOR DEPOT SUSPENSION, LEUPROLIDE ACETATE
 LEVALBUTEROL HYDROCHLORIDE, LEVALBUTEROL HYDROCHLORIDE
 LEVETIRACETAM, LEVETIRACETAM
 LEVETIRACETAM IN SODIUM CHLORIDE, LEVETIRACETAM
 LEVO-T, LEVOTHYROXINE SODIUM **
 LEVOBUNOLOL HYDROCHLORIDE, LEVOBUNOLOL HYDROCHLORIDE
 LEVOCARNITINE, LEVOCARNITINE
 LEVOCARNITINE SF, LEVOCARNITINE
 LEVOCETIRIZINE DIHYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE (OTC)
 LEVOCETIRIZINE DIHYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE
 LEVOCETIRIZINE HYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE
 LEVOFLOXACIN, LEVOFLOXACIN
 LEVOFLOXACIN IN DEXTROSE 5% IN PLASTIC CONTAINER, LEVOFLOXACIN
 LEVOLET, LEVOTHYROXINE SODIUM **
 LEVOLEUCOVORIN CALCIUM, LEVOLEUCOVORIN CALCIUM
 LEVOMILNACIPRAN HYDROCHLORIDE, LEVOMILNACIPRAN HYDROCHLORIDE
 LEVONEST, ETHINYL ESTRADIOL
 LEVONORGESTREL, LEVONORGESTREL (OTC)
 LEVONORGESTREL AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
 LEVONORGESTREL AND ETHINYL ESTRADIOL AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
 LEVONORGESTREL AND ETHINYL ESTRADIOL AND FERROUS FUMARATE, ETHINYL ESTRADIOL
 LEVOPHED, NOREPINEPHRINE BITARTRATE
 LEVORA 0.15/30-28, ETHINYL ESTRADIOL
 LEVORPHANOL TARTRATE, LEVORPHANOL TARTRATE
 LEVOTHYROXINE SODIUM, LEVOTHYROXINE SODIUM
 LEVOTHYROXINE SODIUM, LEVOTHYROXINE SODIUM **
 LEVOXYL, LEVOTHYROXINE SODIUM **
 LEVULAN, AMINOLEVULINIC ACID HYDROCHLORIDE
 LEXAPRO, ESCITALOPRAM OXALATE
 LEXETTE, HALOBETASOL PROPIONATE
 LEXISCAN, REGADENOSON
 LIALDA, MESALAMINE
 LIBERVANT, DIAZEPAM
 LIBRAX, CHLORDIAZEPOXIDE HYDROCHLORIDE
 LIBRIUM, CHLORDIAZEPOXIDE HYDROCHLORIDE
 LICART, DICLOFENAC EPOLAMINE
 LIDOCAINE, LIDOCAINE
 LIDOCAINE AND PRILOCAINE, LIDOCAINE
 LIDOCAINE HYDROCHLORIDE, LIDOCAINE HYDROCHLORIDE
 LIDOCAINE HYDROCHLORIDE 0.2% AND DEXTROSE 5% IN PLASTIC CONTAINER, LIDOCAINE HYDROCHLORIDE
 LIDOCAINE HYDROCHLORIDE 0.4% AND DEXTROSE 5% IN PLASTIC CONTAINER, LIDOCAINE HYDROCHLORIDE
 LIDOCAINE HYDROCHLORIDE 0.8% AND DEXTROSE 5% IN PLASTIC CONTAINER, LIDOCAINE HYDROCHLORIDE
 LIDOCAINE HYDROCHLORIDE 5% AND DEXTROSE 7.5%, LIDOCAINE HYDROCHLORIDE

APPENDIX A - PRODUCT NAME INDEX

** L **

LIDOCAINE HYDROCHLORIDE AND EPINEPHRINE, EPINEPHRINE
LIDOCAINE HYDROCHLORIDE IN PLASTIC CONTAINER, LIDOCAINE HYDROCHLORIDE
LIDOCAINE HYDROCHLORIDE PRESERVATIVE FREE, LIDOCAINE HYDROCHLORIDE
LIDOCAINE HYDROCHLORIDE PRESERVATIVE FREE IN PLASTIC CONTAINER, LIDOCAINE HYDROCHLORIDE
LIDOCAINE HYDROCHLORIDE VISCOUS, LIDOCAINE HYDROCHLORIDE
LIDOCAINE VISCOUS, LIDOCAINE HYDROCHLORIDE
LIDODERM, LIDOCAINE
LIGNOSPAN FORTE, EPINEPHRINE BITARTRATE
LIGNOSPAN STANDARD, EPINEPHRINE BITARTRATE
LIKMEZ, METRONIDAZOLE
LILETTA, LEVONORGESTREL
LINAGLIPTIN, LINAGLIPTIN
LINAGLIPTIN AND METFORMIN HYDROCHLORIDE, LINAGLIPTIN
LINCOCIN, LINCOMYCIN HYDROCHLORIDE
LINCOMYCIN HYDROCHLORIDE, LINCOMYCIN HYDROCHLORIDE
LINEZOLID, LINEZOLID
LINEZOLID IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, LINEZOLID
LINZESS, LINACLOTIDE
LIORESAL, BACLOFEN
LIOTHYRONINE SODIUM, LIOTHYRONINE SODIUM
LIPIODOL, ETHIODIZED OIL
LIPITOR, ATORVASTATIN CALCIUM
LIPOFEN, FENOFIBRATE
LIQUID E-Z-PAQUE, BARIUM SULFATE
LIRAGLUTIDE, LIRAGLUTIDE
LISDEXAMFETAMINE DIMESYLATE, LISDEXAMFETAMINE DIMESYLATE
LISINOPRIL, LISINOPRIL
LISINOPRIL AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
LITFULO, RITLECITINIB TOSYLATE
LITHIUM CARBONATE, LITHIUM CARBONATE
LITHIUM CITRATE, LITHIUM CITRATE
LITHOBID, LITHIUM CARBONATE
LITHOSTAT, ACETOHYDROXAMIC ACID
LIVALO, PITAVASTATIN CALCIUM
LIVDELZI, SELADELPAR LYSINE
LIVMARLI, MARALIXIBAT CHLORIDE
LIVTENCITY, MARIBAVIR
LO LOESTRIN FE, ETHINYL ESTRADIOL
LO SIMPESE, ETHINYL ESTRADIOL
LO-MALMOREDE, ETHINYL ESTRADIOL
LO-ZUMANDIMINE, DROSPIRENONE
LOCAMETZ, GALLIUM GA-68 GOZETOTIDE
LOCHOLEST, CHOLESTYRAMINE
LOCHOLEST LIGHT, CHOLESTYRAMINE
LOCOID, HYDROCORTISONE BUTYRATE
LOCOID LIPOCREAM, HYDROCORTISONE BUTYRATE
LODOCO, COLCHICINE
LODOSYN, CARBIDOPA
LOFEXIDINE HYDROCHLORIDE, LOFEXIDINE HYDROCHLORIDE
LOGILIA, ULIPRISTAL ACETATE
LOKELMA, SODIUM ZIRCONIUM CYCLOSILICATE
LOMAIRA, PHENTERMINE HYDROCHLORIDE
LOMOTIL, ATROPINE SULFATE
LONSURF, TIPIRACIL HYDROCHLORIDE
LOPERAMIDE HYDROCHLORIDE, LOPERAMIDE HYDROCHLORIDE (OTC)
LOPERAMIDE HYDROCHLORIDE, LOPERAMIDE HYDROCHLORIDE
LOPERAMIDE HYDROCHLORIDE AND SIMETHICONE, LOPERAMIDE HYDROCHLORIDE (OTC)
LOPID, GEMFIBROZIL
LOPINAVIR AND RITONAVIR, LOPINAVIR
LOPRESSOR, METOPROLOL TARTRATE
LOPRESSOR HCT, HYDROCHLOROTHIAZIDE
LOPROX, CICLOPIROX
LOPURIN, ALLOPURINOL
LORATADINE, LORATADINE (OTC)
LORATADINE AND PSEUDOEPHEDRINE SULFATE, LORATADINE (OTC)

APPENDIX A - PRODUCT NAME INDEX

** L **

LORATADINE REDIDOSE, LORATADINE (OTC)
 LORAZEPAM, LORAZEPAM
 LORAZEPAM INTENSOL, LORAZEPAM
 LORBRENA, LORLATINIB
 LOREEV XR, LORAZEPAM
 LORYNA, DROSPIRENONE
 LOSARTAN POTASSIUM, LOSARTAN POTASSIUM
 LOSARTAN POTASSIUM AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 LOSEASONIQUE, ETHINYL ESTRADIOL
 LOTEMAX, LOTEPRDNOL ETABONATE
 LOTEMAX SM, LOTEPRDNOL ETABONATE
 LOTENSIN, BENAZEPRIL HYDROCHLORIDE
 LOTENSIN HCT, BENAZEPRIL HYDROCHLORIDE
 LOTEPRDNOL ETABONATE, LOTEPRDNOL ETABONATE
 LOTREL, AMLODIPINE BESYLATE
 LOTRIMIN ULTRA, BUTENAFINE HYDROCHLORIDE (OTC)
 LOTRONEX, ALOSETRON HYDROCHLORIDE
 LOVASTATIN, LOVASTATIN
 LOVAZA, OMEGA-3-ACID ETHYL ESTERS
 LOVENOX, ENOXAPARIN SODIUM
 LOVENOX (PRESERVATIVE FREE), ENOXAPARIN SODIUM
 LOW-OGESTREL-28, ETHINYL ESTRADIOL
 LOXAPINE SUCCINATE, LOXAPINE SUCCINATE
 LUBIPROSTONE, LUBIPROSTONE
 LUCEMYRA, LOFEXIDINE HYDROCHLORIDE
 LUMAKRAS, SOTORASIB
 LUMASON, SULFUR HEXAFLUORIDE LIPID-TYPE A MICROSPHERES
 LUMI-SPORYN, BACITRACIN ZINC
 LUMIFY, BRIMONIDINE TARTRATE (OTC)
 LUMIFY PRESERVATIVE FREE, BRIMONIDINE TARTRATE (OTC)
 LUMIGAN, BIMATOPROST
 LUMISIGHT, PEGULICIANINE ACETATE
 LUMRYZ, SODIUM OXYBATE
 LUNESTA, ESZOPICLONE
 LUPKYNIS, VOCLOSPORIN
 LUPRON DEPOT, LEUPROLIDE ACETATE
 LUPRON DEPOT-PED KIT, LEUPROLIDE ACETATE
 LURASIDONE HYDROCHLORIDE, LURASIDONE HYDROCHLORIDE
 LUTATHERA, LUTETIUM LU 177 DOTATATE
 LUVOX, FLUVOXAMINE MALEATE
 LUZU, LULICONAZOLE
 LYBALVI, OLANZAPINE
 LYMPHOSEEK KIT, TECHNETIUM TC-99M TILMANOCEPT
 LYNPARZA, OLAPARIB
 LYPQOZET, ATORVASTATIN CALCIUM
 LYRICA, PREGABALIN
 LYRICA CR, PREGABALIN
 LYSODREN, MITOTANE
 LYSTEDA, TRANEXAMIC ACID
 LYTGABI, FUTIBATINIB
 LYVISPAH, BACLOFEN

** M **

M-ZOLE 3 COMBINATION PACK, MICONAZOLE NITRATE (OTC)
 MACROBID, NITROFURANTOIN
 MACRODANTIN, NITROFURANTOIN, MACROCRYSTALLINE
 MAGNESIUM SULFATE, MAGNESIUM SULFATE
 MAGNESIUM SULFATE IN DEXTROSE 5% IN PLASTIC CONTAINER, MAGNESIUM SULFATE
 MAGNESIUM SULFATE IN PLASTIC CONTAINER, MAGNESIUM SULFATE
 MALARONE, ATOVAQUONE
 MALARONE PEDIATRIC, ATOVAQUONE
 MALATHION, MALATHION
 MALMOREDE, ETHINYL ESTRADIOL
 MANGANESE CHLORIDE IN PLASTIC CONTAINER, MANGANESE CHLORIDE
 MANNITOL 10% IN PLASTIC CONTAINER, MANNITOL

APPENDIX A - PRODUCT NAME INDEX

** M **

MANNITOL 15% IN PLASTIC CONTAINER, MANNITOL
MANNITOL 20% IN PLASTIC CONTAINER, MANNITOL
MANNITOL 25%, MANNITOL
MANNITOL 5% IN PLASTIC CONTAINER, MANNITOL
MARAVIROC, MARAVIROC
MARCINE, BUPIVACAINE HYDROCHLORIDE
MARCINE HYDROCHLORIDE, BUPIVACAINE HYDROCHLORIDE
MARCINE HYDROCHLORIDE PRESERVATIVE FREE, BUPIVACAINE HYDROCHLORIDE
MARCINE HYDROCHLORIDE W/ EPINEPHRINE, BUPIVACAINE HYDROCHLORIDE
MARCINE HYDROCHLORIDE W/ EPINEPHRINE PRESERVATIVE FREE, BUPIVACAINE HYDROCHLORIDE
MARINOL, DRONABINOL
MARLISSA, ETHINYL ESTRADIOL
MARPLAN, ISOCARBOXAZID
MATULANE, PROCARBAZINE HYDROCHLORIDE
MAVENCLAD, CLADRIBINE
MAVYRET, GLECAPREVIR
MAXALT, RIZATRIPTAN BENZOATE
MAXALT-MLT, RIZATRIPTAN BENZOATE
MAXIDEX, DEXAMETHASONE
MAXITROL, DEXAMETHASONE
MAXZIDE, HYDROCHLOROTHIAZIDE
MAXZIDE-25, HYDROCHLOROTHIAZIDE
MAYZENT, SIPONIMOD
MD-GASTROVIEW, DIATRIZOATE MEGLUMINE
MECAMYLAMINE HYDROCHLORIDE, MECAMYLAMINE HYDROCHLORIDE
MECLIZINE HYDROCHLORIDE, MECLIZINE HYDROCHLORIDE
MECLOFENAMATE SODIUM, MECLOFENAMATE SODIUM
MEDROL, METHYLPREDNISOLONE
MEDROXYPROGESTERONE ACETATE, MEDROXYPROGESTERONE ACETATE
MEFENAMIC ACID, MEFENAMIC ACID
MEFLOQUINE HYDROCHLORIDE, MEFLOQUINE HYDROCHLORIDE
MEGACE ES, MEGESTROL ACETATE
MEGESTROL ACETATE, MEGESTROL ACETATE
MEKINIST, TRAMETINIB DIMETHYL SULFOXIDE
MEKTOVI, BINIMETINIB
MELAMISA, DROSPIRENONE
MELOXICAM, MELOXICAM
MELPHALAN HYDROCHLORIDE, MELPHALAN HYDROCHLORIDE
MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
MEMANTINE HYDROCHLORIDE AND DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
MEN'S ROGAINE, MINOXIDIL (OTC)
MENOSTAR, ESTRADIOL
MEPERIDINE HYDROCHLORIDE, MEPERIDINE HYDROCHLORIDE
MEPERIDINE HYDROCHLORIDE PRESERVATIVE FREE, MEPERIDINE HYDROCHLORIDE
MEPROBAMATE, MEPROBAMATE
MEPRON, ATOVAQUONE
MERCAPTOPYRINE, MERCAPTOPYRINE
MEROPENEM, MEROPENEM
MEROPENEM AND SODIUM CHLORIDE IN DUPLEX CONTAINER, MEROPENEM
MERREM IV, MEROPENEM
MERZEE, ETHINYL ESTRADIOL
MESALAMINE, MESALAMINE
MESNA, MESNA
MESNEX, MESNA
MESTINON, PYRIDOSTIGMINE BROMIDE
METADATE CD, METHYLPHENIDATE HYDROCHLORIDE
METAPROTERENOL SULFATE, METAPROTERENOL SULFATE
METARAMINOL BITARTRATE, METARAMINOL BITARTRATE
METASTRON, STRONTIUM CHLORIDE SR-89
METAXALONE, METAXALONE
METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
METFORMIN HYDROCHLORIDE; SAXAGLIPTIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
METHADONE HYDROCHLORIDE, METHADONE HYDROCHLORIDE
METHADONE HYDROCHLORIDE INTENSOL, METHADONE HYDROCHLORIDE
METHADOSE, METHADONE HYDROCHLORIDE

APPENDIX A - PRODUCT NAME INDEX

** M **

METHAMPHETAMINE HYDROCHLORIDE, METHAMPHETAMINE HYDROCHLORIDE
 METHAZOLAMIDE, METHAZOLAMIDE
 METHENAMINE HIPPURATE, METHENAMINE HIPPURATE
 METHIMAZOLE, METHIMAZOLE
 METHOCARBAMOL, METHOCARBAMOL
 METHOHEXITAL SODIUM, METHOHEXITAL SODIUM
 METHOTREXATE PRESERVATIVE FREE, METHOTREXATE SODIUM
 METHOTREXATE SODIUM, METHOTREXATE SODIUM
 METHOTREXATE SODIUM PRESERVATIVE FREE, METHOTREXATE SODIUM
 METHOXSALEN, METHOXSALEN
 METHSCOPOLAMINE BROMIDE, METHSCOPOLAMINE BROMIDE
 METHSUXIMIDE, METHSUXIMIDE
 METHYLDOPA, METHYLDOPA
 METHYLENE BLUE, METHYLENE BLUE
 METHYLERGONOVINE MALEATE, METHYLERGONOVINE MALEATE
 METHYLIN, METHYLPHENIDATE HYDROCHLORIDE
 METHYLIN ER, METHYLPHENIDATE HYDROCHLORIDE
 METHYLNALTREXONE BROMIDE, METHYLNALTREXONE BROMIDE
 METHYLPHENIDATE, METHYLPHENIDATE
 METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
 METHYLPREDNISOLONE, METHYLPREDNISOLONE
 METHYLPREDNISOLONE ACETATE, METHYLPREDNISOLONE ACETATE
 METHYLPREDNISOLONE SODIUM SUCCINATE, METHYLPREDNISOLONE SODIUM SUCCINATE
 METHYLTESTOSTERONE, METHYLTESTOSTERONE
 METOCLOPRAMIDE, METOCLOPRAMIDE HYDROCHLORIDE
 METOCLOPRAMIDE HYDROCHLORIDE, METOCLOPRAMIDE HYDROCHLORIDE
 METOLAZONE, METOLAZONE
 METOPIRONE, METYRAPONE
 METOPROLOL SUCCINATE, METOPROLOL SUCCINATE
 METOPROLOL TARTRATE, METOPROLOL TARTRATE
 METOPROLOL TARTRATE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 METRO I.V. IN PLASTIC CONTAINER, METRONIDAZOLE
 METROCREAM, METRONIDAZOLE
 METROGEL, METRONIDAZOLE
 METROGEL-VAGINAL, METRONIDAZOLE
 METROLOTION, METRONIDAZOLE
 METRONIDAZOLE, METRONIDAZOLE
 METRONIDAZOLE IN PLASTIC CONTAINER, METRONIDAZOLE
 METYROSINE, METYROSINE
 MEXILETINE HYDROCHLORIDE, MEXILETINE HYDROCHLORIDE
 MIACALCIN, CALCITONIN SALMON
 MIBELAS 24 FE, ETHINYL ESTRADIOL
 MICA FUNGIN, MICA FUNGIN SODIUM
 MICA FUNGIN IN SODIUM CHLORIDE 0.9%, MICA FUNGIN SODIUM
 MICA FUNGIN SODIUM, MICA FUNGIN SODIUM
 MICARDIS, TELMISARTAN
 MICARDIS HCT, HYDROCHLOROTHIAZIDE
 MICONAZOLE 3, MICONAZOLE NITRATE (OTC)
 MICONAZOLE 3 COMBINATION PACK, MICONAZOLE NITRATE (OTC)
 MICONAZOLE 7, MICONAZOLE NITRATE (OTC)
 MICONAZOLE NITRATE, MICONAZOLE NITRATE (OTC)
 MICONAZOLE NITRATE, MICONAZOLE NITRATE
 MICONAZOLE NITRATE COMBINATION PACK, MICONAZOLE NITRATE (OTC)
 MICORT-HC, HYDROCORTISONE ACETATE
 MICROZIDE, HYDROCHLOROTHIAZIDE
 MIDAMOR, AMILORIDE HYDROCHLORIDE
 MIDAZOLAM HYDROCHLORIDE, MIDAZOLAM HYDROCHLORIDE
 MIDAZOLAM HYDROCHLORIDE (AUTOINJECTOR), MIDAZOLAM HYDROCHLORIDE
 MIDAZOLAM HYDROCHLORIDE PRESERVATIVE FREE, MIDAZOLAM HYDROCHLORIDE
 MIDAZOLAM IN 0.9% SODIUM CHLORIDE, MIDAZOLAM
 MIDODRINE HYDROCHLORIDE, MIDODRINE HYDROCHLORIDE
 MIDOL LIQUID GELS, IBUPROFEN (OTC)
 MIDOSTAURIN, MIDOSTAURIN
 MIDOZALAM HYDROCHLORIDE, MIDAZOLAM HYDROCHLORIDE
 MIEBO, PERFLUOROHEXYLOCTANE

APPENDIX A - PRODUCT NAME INDEX

** M **

MIFEPREX, MIFEPRISTONE
MIFEPRISTONE, MIFEPRISTONE
MIGERGOT, CAFFEINE
MIGLITOL, MIGLITOL
MIGLUSTAT, MIGLUSTAT
MIGRANAL, DIHYDROERGOTAMINE MESYLATE
MILI, ETHINYL ESTRADIOL
MILNACIPRAN HYDROCHLORIDE, MILNACIPRAN HYDROCHLORIDE
MILRINONE LACTATE, MILRINONE LACTATE
MILRINONE LACTATE IN DEXTROSE 5%, MILRINONE LACTATE
MILRINONE LACTATE IN DEXTROSE 5% IN PLASTIC CONTAINER, MILRINONE LACTATE
MILRINONE LACTATE IN PLASTIC CONTAINER, MILRINONE LACTATE
MINIPRESS, PRAZOSIN HYDROCHLORIDE
MINIVELLE, ESTRADIOL
MINOCIN, MINOCYCLINE HYDROCHLORIDE
MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE
MINOXIDIL, MINOXIDIL (OTC)
MINOXIDIL, MINOXIDIL
MINOXIDIL (FOR MEN), MINOXIDIL (OTC)
MINOXIDIL (FOR WOMEN), MINOXIDIL (OTC)
MINOXIDIL EXTRA STRENGTH (FOR MEN), MINOXIDIL (OTC)
MINZOYA, ETHINYL ESTRADIOL
MIOCHOL-E, ACETYLCHOLINE CHLORIDE
MIOSTAT, CARBACHOL
MIPLYFFA, ARIMOCLOMOL CITRATE
MIRABEGRON, MIRABEGRON
MIRALAX, POLYETHYLENE GLYCOL 3350 (OTC)
MIRENA, LEVONORGESTREL
MIRTAZAPINE, MIRTAZAPINE
MIRVASO, BRIMONIDINE TARTRATE
MISOPROSTOL, MISOPROSTOL
MITIGARE, COLCHICINE
MITIGO, MORPHINE SULFATE
MITOMYCIN, MITOMYCIN
MITOSOL, MITOMYCIN
MITOXANTRONE HYDROCHLORIDE, MITOXANTRONE HYDROCHLORIDE
MODAFINIL, MODAFINIL
MOEXIPRIL HYDROCHLORIDE, MOEXIPRIL HYDROCHLORIDE
MOEXIPRIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
MOLINDONE HYDROCHLORIDE, MOLINDONE HYDROCHLORIDE
MOMETASONE FUROATE, MOMETASONE FUROATE (OTC)
MOMETASONE FUROATE, MOMETASONE FUROATE
MONISTAT 1 COMBINATION PACK, MICONAZOLE NITRATE (OTC)
MONISTAT 3, MICONAZOLE NITRATE (OTC)
MONISTAT 3, MICONAZOLE NITRATE
MONISTAT 3 COMBINATION PACK, MICONAZOLE NITRATE (OTC)
MONISTAT 3 COMBINATION PACK (PREFILLED), MICONAZOLE NITRATE (OTC)
MONISTAT 7, MICONAZOLE NITRATE (OTC)
MONISTAT 7 COMBINATION PACK, MICONAZOLE NITRATE (OTC)
MONO-LINYAH, ETHINYL ESTRADIOL
MONOBASIC SODIUM PHOSPHATE AND DIBASIC SODIUM PHOSPHATE, SODIUM PHOSPHATE, DIBASIC,
MONODOX, DOXYCYCLINE
MONOFERRIC, FERRIC DERISOMALTOSE
MONOKET, ISOSORBIDE MONONITRATE
MONTELUKAST SODIUM, MONTELUKAST SODIUM
MORPHINE SULFATE, MORPHINE SULFATE
MOTTEGRITY, PRUCALOPRIDE SUCCINATE
MOTOFEN, ATROPINE SULFATE
MOTPOLY XR, LACOSAMIDE
MOTRIN IB, IBUPROFEN (OTC)
MOUNJARO, TIRZEPATIDE
MOUNJARO (AUTOINJECTOR), TIRZEPATIDE
MOVANTIK, NALOXEGOL OXALATE
MOVIPREP, ASCORBIC ACID
MOXIDECTIN, MOXIDECTIN

APPENDIX A - PRODUCT NAME INDEX**** M ****

MOXIFLOXACIN HYDROCHLORIDE, MOXIFLOXACIN HYDROCHLORIDE
 MOXIFLOXACIN HYDROCHLORIDE IN SODIUM CHLORIDE 0.8% IN PLASTIC CONTAINER, MOXIFLOXACIN
 MOZOBIL, PLERIXAFOR
 MPI INDIUM DTPA IN 111, INDIUM IN-111 PENTETATE DISODIUM
 MS CONTIN, MORPHINE SULFATE
 MUCINEX, GUAIFENESIN (OTC)
 MUCINEX D, GUAIFENESIN (OTC)
 MUCINEX DM, DEXTROMETHORPHAN HYDROBROMIDE (OTC)
 MULPLETA, LUSUTROMBOPAG
 MULTAQ, DRONEDARONE HYDROCHLORIDE
 MULTIHANCE, GADOBENATE DIMEGLUMINE
 MULTIHANCE MULTIPACK, GADOBENATE DIMEGLUMINE
 MULTIPLE ELECTROLYTES INJECTION TYPE 1 USP PH 5.5, MAGNESIUM CHLORIDE
 MULTIPLE ELECTROLYTES INJECTION TYPE 1 USP PH 7.4, MAGNESIUM CHLORIDE
 MULTRYS, CUPRIC SULFATE
 MUPIROCIN, MUPIROCIN
 MUPIROCIN, MUPIROCIN CALCIUM
 MYAMBUTOL, ETHAMBUTOL HYDROCHLORIDE
 MYCAMINE, MICAFUNGIN SODIUM
 MYCAPSSA, OCTREOTIDE ACETATE
 MYCOBUTIN, RIFABUTIN
 MYCOPHENOLATE MOFETIL, MYCOPHENOLATE MOFETIL
 MYCOPHENOLATE MOFETIL HYDROCHLORIDE, MYCOPHENOLATE MOFETIL HYDROCHLORIDE
 MYCOPHENOLIC SODIUM, MYCOPHENOLIC SODIUM
 MYDAYIS, AMPHETAMINE ASPARTATE
 MYDRIACYL, TROPICAMIDE
 MYFEMBREE, ESTRADIOL
 MYFORTIC, MYCOPHENOLIC SODIUM
 MYHIBBIN, MYCOPHENOLATE MOFETIL
 MYLERAN, BUSULFAN
 MYORISAN, ISOTRETINOIN
 MYOVIEW, TECHNETIUM TC-99M TETROFOSMIN KIT
 MYOVIEW 30ML, TECHNETIUM TC-99M TETROFOSMIN KIT
 MYRBETRIQ, MIRABEGRON
 MYRBETRIQ GRANULES, MIRABEGRON
 MYSOLINE, PRIMIDONE
 MYTESI, CROFELEMER
 MYZILRA, ETHINYL ESTRADIOL

**** N ****

NABUMETONE, NABUMETONE
 NADOLOL, NADOLOL
 NAFCILLIN SODIUM, NAFCILLIN SODIUM
 NAFTIFINE HYDROCHLORIDE, NAFTIFINE HYDROCHLORIDE
 NAFTIN, NAFTIFINE HYDROCHLORIDE
 NALBUPHINE HYDROCHLORIDE, NALBUPHINE HYDROCHLORIDE
 NALLPEN IN PLASTIC CONTAINER, NAFCILLIN SODIUM
 NALMEFENE HYDROCHLORIDE, NALMEFENE HYDROCHLORIDE
 NALOXONE, NALOXONE HYDROCHLORIDE
 NALOXONE HYDROCHLORIDE, NALOXONE HYDROCHLORIDE (OTC)
 NALOXONE HYDROCHLORIDE, NALOXONE HYDROCHLORIDE
 NALOXONE HYDROCHLORIDE AND PENTAZOCINE HYDROCHLORIDE, NALOXONE HYDROCHLORIDE
 NALTREXONE HYDROCHLORIDE, NALTREXONE HYDROCHLORIDE
 NAMENDA, MEMANTINE HYDROCHLORIDE
 NAMZARIC, DONEPEZIL HYDROCHLORIDE
 NAPHAZOLINE HYDROCHLORIDE AND PHENIRAMINE MALEATE, NAPHAZOLINE HYDROCHLORIDE (OTC)
 NAPHCN-A, NAPHAZOLINE HYDROCHLORIDE (OTC)
 NAPRELAN, NAPROXEN SODIUM
 NAPROSYN, NAPROXEN
 NAPROXEN, NAPROXEN
 NAPROXEN AND ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM
 NAPROXEN SODIUM, NAPROXEN SODIUM (OTC)
 NAPROXEN SODIUM, NAPROXEN SODIUM
 NAPROXEN SODIUM AND DIPHENHYDRAMINE HYDROCHLORIDE, DIPHENHYDRAMINE HYDROCHLORIDE (OTC)
 NAPROXEN SODIUM AND PSEUDOEPHEDRINE HYDROCHLORIDE, NAPROXEN SODIUM (OTC)

APPENDIX A - PRODUCT NAME INDEX

** N **

NARATRIPTAN, NARATRIPTAN HYDROCHLORIDE
NARCAN, NALOXONE HYDROCHLORIDE (OTC)
NARDIL, PHENELZINE SULFATE
NAROPIN, ROPIVACAINE HYDROCHLORIDE
NASACORT ALLERGY 24 HOUR, TRIAMCINOLONE ACETONIDE (OTC)
NASONEX 24HR ALLERGY, MOMETASONE FUROATE (OTC)
NATACYN, NATAMYCIN
NATAZIA, DIENOGEST
NATEGLINIDE, NATEGLINIDE
NATESTO, TESTOSTERONE
NATROBA, SPINOSAD
NAYZILAM, MIDAZOLAM
NEBIVOLOL HYDROCHLORIDE, NEBIVOLOL HYDROCHLORIDE
NEBUPENT, PENTAMIDINE ISETHIONATE
NEFAZODONE HYDROCHLORIDE, NEFAZODONE HYDROCHLORIDE
NEFFY, EPINEPHRINE
NELARABINE, NELARABINE
NEMBUTAL SODIUM, PENTOBARBITAL SODIUM
NEO-SYNALAR, FLUOCINOLONE ACETONIDE
NEOMYCIN AND POLYMYXIN B SULFATE, NEOMYCIN SULFATE
NEOMYCIN AND POLYMYXIN B SULFATES AND BACITRACIN ZINC, BACITRACIN ZINC
NEOMYCIN AND POLYMYXIN B SULFATES AND DEXAMETHASONE, DEXAMETHASONE
NEOMYCIN AND POLYMYXIN B SULFATES AND GRAMICIDIN, GRAMICIDIN
NEOMYCIN AND POLYMYXIN B SULFATES AND HYDROCORTISONE, HYDROCORTISONE
NEOMYCIN AND POLYMYXIN B SULFATES, BACITRACIN ZINC AND HYDROCORTISONE, BACITRACIN ZINC
NEOMYCIN SULFATE, NEOMYCIN SULFATE
NEOPROFEN, IBUPROFEN LYSINE
NEORAL, CYCLOSPORINE
NEOSPORIN, GRAMICIDIN
NEOSPORIN G.U. IRRIGANT, NEOMYCIN SULFATE
NEOSTIGMINE METHYLSULFATE, NEOSTIGMINE METHYLSULFATE
NEOSTIGMINE METHYLSULFATE, NEOSTIGMINE METHYLSULFATE
NEPHROSCAN, TECHNETIUM TC-99M SUCCIMER
NERLYNX, NERATINIB MALEATE
NESACAINE, CHLOROPROCAINE HYDROCHLORIDE
NESACAINE-MPF, CHLOROPROCAINE HYDROCHLORIDE
NESINA, ALOGLIPTIN BENZOATE
NETSPOT, GALLIUM DOTATATE GA-68
NEUPRO, ROTIGOTINE
NEURACEQ, FLORBETABEN F-18
NEUROLITE, TECHNETIUM TC-99M BICISATE KIT
NEURONTIN, GABAPENTIN
NEVANAC, NEPAFENAC
NEVIRAPINE, NEVIRAPINE
NEXAVAR, SORAFENIB TOSYLATE
NEXESTA FE, ETHINYL ESTRADIOL
NEXICLON XR, CLONIDINE
NEXIUM, ESOMEPRAZOLE MAGNESIUM
NEXIUM 24HR, ESOMEPRAZOLE MAGNESIUM (OTC)
NEXLETOL, BEMPEDOIC ACID
NEXLIZET, BEMPEDOIC ACID
NEXPLANON, ETONOGESTREL
NEXTERONE, AMIODARONE HYDROCHLORIDE
NEXTSTELLIS, DROSPIRENONE
NIACIN, NIACIN
NIACOR, NIACIN
NICARDIPINE HYDROCHLORIDE, NICARDIPINE HYDROCHLORIDE
NICARDIPINE HYDROCHLORIDE IN 0.83% SODIUM CHLORIDE, NICARDIPINE HYDROCHLORIDE
NICARDIPINE HYDROCHLORIDE IN 0.86% SODIUM CHLORIDE, NICARDIPINE HYDROCHLORIDE
NICARDIPINE HYDROCHLORIDE IN 0.9% SODIUM CHLORIDE, NICARDIPINE HYDROCHLORIDE
NICODERM CQ, NICOTINE (OTC)
NICORETTE, NICOTINE POLACRILEX (OTC)
NICORETTE (MINT), NICOTINE POLACRILEX (OTC)
NICOTINE, NICOTINE (OTC)
NICOTINE POLACRILEX, NICOTINE POLACRILEX (OTC)

APPENDIX A - PRODUCT NAME INDEX

** N **

NICOTROL, NICOTINE
 NIFEDIPINE, NIFEDIPINE
 NIKKI, DROSPIRENONE
 NILANDRON, NILUTAMIDE
 NILUTAMIDE, NILUTAMIDE
 NIMODIPINE, NIMODIPINE
 NINLARO, IXAZOMIB CITRATE
 NIPENT, PENTOSTATIN
 NIPRIDE RTU IN SODIUM CHLORIDE 0.9%, SODIUM NITROPRUSSIDE
 NISOLDIPINE, NISOLDIPINE
 NITAZOXANIDE, NITAZOXANIDE
 NITHIODOLE, SODIUM NITRITE
 NITISINONE, NITISINONE
 NITRO-DUR, NITROGLYCERIN
 NITROFURANTOIN, NITROFURANTOIN
 NITROFURANTOIN, NITROFURANTOIN, MACROCRYSTALLINE
 NITROFURANTOIN (MONOHYDRATE/MACROCRYSTALS), NITROFURANTOIN
 NITROGLYCERIN, NITROGLYCERIN
 NITROGLYCERIN IN DEXTROSE 5%, NITROGLYCERIN
 NITROLINGUAL PUMPSPRAY, NITROGLYCERIN
 NITROMIST, NITROGLYCERIN
 NITROSTAT, NITROGLYCERIN
 NITYR, NITISINONE
 NIX, PERMETHRIN (OTC)
 NIZATIDINE, NIZATIDINE
 NIZORAL ANTI-DANDRUFF, KETOCONAZOLE (OTC)
 NOR-QD, NORETHINDRONE
 NOREPINEPHRINE BITARTRATE, NOREPINEPHRINE BITARTRATE
 NOREPINEPHRINE BITARTRATE IN 0.9% SODIUM CHLORIDE, NOREPINEPHRINE BITARTRATE
 NOREPINEPHRINE BITARTRATE IN 5% DEXTROSE, NOREPINEPHRINE BITARTRATE
 NORETHINDRONE, NORETHINDRONE
 NORETHINDRONE ACETATE, NORETHINDRONE ACETATE
 NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
 NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL AND FERROUS FUMARATE, ETHINYL ESTRADIOL
 NORETHINDRONE AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
 NORETHINDRONE AND ETHINYL ESTRADIOL (10/11), ETHINYL ESTRADIOL
 NORETHINDRONE AND ETHINYL ESTRADIOL AND FERROUS FUMARATE, ETHINYL ESTRADIOL
 NORGESTIMATE AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
 NORITATE, METRONIDAZOLE
 NORLIQVA, AMLODIPINE BESYLATE
 NORMOCARB HF 25, MAGNESIUM CHLORIDE
 NORMOCARB HF 35, MAGNESIUM CHLORIDE
 NORMOSOL-M AND DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
 NORMOSOL-R AND DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
 NORMOSOL-R IN PLASTIC CONTAINER, MAGNESIUM CHLORIDE
 NORPACE, DISOPYRAMIDE PHOSPHATE
 NORPACE CR, DISOPYRAMIDE PHOSPHATE
 NORPRAMIN, DESIPRAMINE HYDROCHLORIDE
 NORTHERA, DROXIDOPA
 NORTREL 0.5/35-28, ETHINYL ESTRADIOL
 NORTREL 1/35-21, ETHINYL ESTRADIOL
 NORTREL 1/35-28, ETHINYL ESTRADIOL
 NORTREL 7/7/7, ETHINYL ESTRADIOL
 NORTRIPTYLINE HYDROCHLORIDE, NORTRIPTYLINE HYDROCHLORIDE
 NORVASC, AMLODIPINE BESYLATE
 NORVIR, RITONAVIR
 NOURIANZ, ISTRADefylline
 NOXAFIL, POSACONAZOLE
 NOXAFIL POWDERMIX KIT, POSACONAZOLE
 NOXIVENT, NITRIC OXIDE
 NUBEQA, DAROLUTAMIDE
 NUCYNТА, TAPENTADOL HYDROCHLORIDE
 NUCYNТА ER, TAPENTADOL HYDROCHLORIDE
 NUEDEXТА, DEXTROMETHORPHAN HYDROBROMIDE
 NULIBRY, FOSDENOPTERIN HYDROBROMIDE

APPENDIX A - PRODUCT NAME INDEX

** N **

NULYTELY, POLYETHYLENE GLYCOL 3350
 NULYTELY-FLAVORED, POLYETHYLENE GLYCOL 3350
 NUMBRINO, COCAINE HYDROCHLORIDE
 NUPLAZID, PIMAVANSERIN TARTRATE
 NURTEC ODT, RIMEGEPANT SULFATE
 NUTRILIPID 20%, SOYBEAN OIL
 NUVARING, ETHINYL ESTRADIOL
 NUVESSA, METRONIDAZOLE
 NUVIGIL, ARMODAFINIL
 NUZYRA, OMADACYCLINE TOSYLATE
 NYLIA 1/35, ETHINYL ESTRADIOL
 NYLIA 7/7/7, ETHINYL ESTRADIOL
 NYMALIZE, NIMODIPINE
 NYSTATIN, NYSTATIN
 NYSTATIN AND TRIAMCINOLONE ACETONIDE, NYSTATIN
 NYSTOP, NYSTATIN

** O **

OCALIVA, OBETICHOLIC ACID
 OCTREOSCAN, INDIUM IN-111 PENTETREOTIDE KIT
 OCTREOTIDE ACETATE, OCTREOTIDE ACETATE
 OCTREOTIDE ACETATE (PRESERVATIVE FREE), OCTREOTIDE ACETATE
 OCUFLOX, OFLOXACIN
 ODEFSEY, EMTRICITABINE
 ODOMZO, SONIDEGIB PHOSPHATE
 OFEV, NINTEDANIB ESYLATE
 OFLOXACIN, OFLOXACIN
 OGEN 5, ESTROPIPATE
 OGSIVEO, NIROGACESTAT HYDROBROMIDE
 OHTUVAYRE, ENSIFENTRINE
 OJEMDA, TOVORAFENIB
 OJJAARA, MOMELOTINIB DIHYDROCHLORIDE
 OLANZAPINE, OLANZAPINE
 OLANZAPINE AND FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 OLINVYK, OLICERIDINE
 OLMESARTAN MEDOXOMIL, OLMESARTAN MEDOXOMIL
 OLMESARTAN MEDOXOMIL AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 OLMESARTAN MEDOXOMIL, AMLODIPINE AND HYDROCHLOROTHIAZIDE, AMLODIPINE BESYLATE
 OLOPATADINE HYDROCHLORIDE, OLOPATADINE HYDROCHLORIDE (OTC)
 OLOPATADINE HYDROCHLORIDE, OLOPATADINE HYDROCHLORIDE
 OLPRUVA, SODIUM PHENYL BUTYRATE
 OLUMIANT, BARICITINIB
 OMEGA-3-ACID ETHYL ESTERS, OMEGA-3-ACID ETHYL ESTERS
 OMEGAVEN, FISH OIL TRIGLYCERIDES
 OMEPRAZOLE, OMEPRAZOLE (OTC)
 OMEPRAZOLE, OMEPRAZOLE
 OMEPRAZOLE AND SODIUM BICARBONATE, OMEPRAZOLE (OTC)
 OMEPRAZOLE AND SODIUM BICARBONATE, OMEPRAZOLE
 OMEPRAZOLE MAGNESIUM, OMEPRAZOLE MAGNESIUM (OTC)
 OMIDRIA, KETOROLAC TROMETHAMINE
 OMNARIS, CICLESONIDE
 OMNIPAQUE 12, IOHEXOL
 OMNIPAQUE 140, IOHEXOL
 OMNIPAQUE 180, IOHEXOL
 OMNIPAQUE 240, IOHEXOL
 OMNIPAQUE 300, IOHEXOL
 OMNIPAQUE 350, IOHEXOL
 OMNIPAQUE 9, IOHEXOL
 OMNIPRED, PREDNISOLONE ACETATE
 OMNISCAN, GADODIAMIDE
 ONDANSETRON, ONDANSETRON
 ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
 ONDANSETRON HYDROCHLORIDE PRESERVATIVE FREE, ONDANSETRON HYDROCHLORIDE
 ONEXTON, BENZOYL PEROXIDE
 ONFI, CLOBAZAM

APPENDIX A - PRODUCT NAME INDEX

** O **

ONGENTYS, OPICAPONE
 ONIVYDE, IRINOTECAN HYDROCHLORIDE
 ONPATTRO, PATISIRAN SODIUM
 ONSURA, ETHINYL ESTRADIOL
 ONTRALFY, TIZANIDINE HYDROCHLORIDE
 ONUREG, AZACITIDINE
 ONYDA XR, CLONIDINE HYDROCHLORIDE
 ONZETRA XSAIL, SUMATRIPTAN SUCCINATE
 OPCICON ONE-STEP, LEVONORGESTREL (OTC)
 OPCON-A, NAPHAZOLINE HYDROCHLORIDE (OTC)
 OPFOLDA, MIGLUSTAT
 OPILL, NORGESTREL (OTC)
 OPIPZA, ARIPIPRAZOLE
 OPSUMIT, MACITENTAN
 OPSYNVI, MACITENTAN
 OPTIRAY 300, IOVERSOL
 OPTIRAY 320, IOVERSOL
 OPTIRAY 350, IOVERSOL
 OPTISON, ALBUMIN HUMAN
 OPVEE, NALMEFENE HYDROCHLORIDE
 OPZELURA, RUXOLITINIB PHOSPHATE
 ORABLOC, ARTICAININE HYDROCHLORIDE
 ORACEA, DOXYCYCLINE
 ORAPRED ODT, PREDNISOLONE SODIUM PHOSPHATE
 ORAQIX, LIDOCAINE
 ORAVERSE, PHENTOLAMINE MESYLATE
 ORAVIG, MICONAZOLE
 ORBACTIV, ORITAVANCIN DIPHOSPHATE
 ORENITRAM, TREPROSTINIL DIOLAMINE
 ORFADIN, NITISINONE
 ORGOVYX, RELUGOLIX
 ORIAHNN (COPACKAGED), ELAGOLIX SODIUM, ESTRADIOL, NORETHINDRONE ACETATE
 ORILISSA, ELAGOLIX SODIUM
 ORKAMBI, IVACAFTOR
 ORLADEYO, BEROTRALSTAT HYDROCHLORIDE
 ORLYNVAH, PROBENECID
 ORMALVI, DICHLORPHENAMIDE
 ORPHENADRINE CITRATE, ORPHENADRINE CITRATE
 ORPHENGESIC, ASPIRIN
 ORPHENGESIC FORTE, ASPIRIN
 ORSERDU, ELACESTRANT HYDROCHLORIDE
 ORVATEN, MIDODRINE HYDROCHLORIDE
 OSELTAMIVIR PHOSPHATE, OSELTAMIVIR PHOSPHATE
 OSENI, ALOGLIPTIN BENZOATE
 OSMITROL 10% IN WATER, MANNITOL
 OSMITROL 10% IN WATER IN PLASTIC CONTAINER, MANNITOL
 OSMITROL 15% IN WATER, MANNITOL
 OSMITROL 15% IN WATER IN PLASTIC CONTAINER, MANNITOL
 OSMITROL 20% IN WATER, MANNITOL
 OSMITROL 20% IN WATER IN PLASTIC CONTAINER, MANNITOL
 OSMITROL 5% IN WATER, MANNITOL
 OSMITROL 5% IN WATER IN PLASTIC CONTAINER, MANNITOL
 OSPEMIFENE, OSPEMIFENE
 OSPHENA, OSPEMIFENE
 OTEZLA, APREMILAST
 OTICAIR, HYDROCORTISONE
 OTOVEL, CIPROFLOXACIN HYDROCHLORIDE
 OTREXUP, METHOTREXATE
 OXACILLIN SODIUM, OXACILLIN SODIUM
 OXALIPLATIN, OXALIPLATIN
 OXAPROZIN, OXAPROZIN
 OXAZEPAM, OXAZEPAM
 OXCARBAZEPINE, OXCARBAZEPINE
 OXICONAZOLE NITRATE, OXICONAZOLE NITRATE
 OXISTAT, OXICONAZOLE NITRATE

APPENDIX A - PRODUCT NAME INDEX

** O **

OXLUMO, LUMASIRAN SODIUM
 OXSORALEN-ULTRA, METHOXSALEN
 OXTELLAR XR, OXCARBAZEPINE
 OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE
 OXYCET, ACETAMINOPHEN
 OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN
 OXYCODONE AND ASPIRIN, ASPIRIN
 OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
 OXYCONTIN, OXYCODONE HYDROCHLORIDE
 OXYMORPHONE HYDROCHLORIDE, OXYMORPHONE HYDROCHLORIDE
 OXYTOCIN, OXYTOCIN
 OXYTROL, OXYBUTYNIN
 OXYTROL FOR WOMEN, OXYBUTYNIN (OTC)
 OZEMPIC, SEMAGLUTIDE
 OZOBAX DS, BACLOFEN
 OZURDEX, DEXAMETHASONE

** P **

PACERONE, AMIODARONE HYDROCHLORIDE
 PACLITAXEL, PACLITAXEL
 PALIPERIDONE, PALIPERIDONE
 PALIPERIDONE PALMITATE, PALIPERIDONE PALMITATE
 PALONOSETRON HYDROCHLORIDE, PALONOSETRON HYDROCHLORIDE
 PAMIDRONATE DISODIUM, PAMIDRONATE DISODIUM
 PANCURONIUM BROMIDE, PANCURONIUM BROMIDE
 PANRETIN, ALITRETINOIN
 PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
 PANTOPRAZOLE SODIUM IN 0.9% SODIUM CHLORIDE, PANTOPRAZOLE SODIUM
 PARAGARD T 380A, COPPER
 PARICALCITOL, PARICALCITOL
 PARLODEL, BROMOCRIPTINE MESYLATE
 PARNATE, TRANYLCPROMINE SULFATE
 PAROMOMYCIN SULFATE, PAROMOMYCIN SULFATE
 PAROXETINE, PAROXETINE HYDROCHLORIDE
 PAROXETINE HYDROCHLORIDE, PAROXETINE HYDROCHLORIDE
 PAROXETINE MESYLATE, PAROXETINE MESYLATE
 PARSABIV, ETELALCETIDE
 PASER, AMINOSALICYLIC ACID
 PATADAY ONCE DAILY RELIEF, OLOPATADINE HYDROCHLORIDE (OTC)
 PATADAY TWICE DAILY RELIEF, OLOPATADINE HYDROCHLORIDE (OTC)
 PAXIL, PAROXETINE HYDROCHLORIDE
 PAXIL CR, PAROXETINE HYDROCHLORIDE
 PAXLOVID (COPACKAGED), NIRMATRELVIR
 PAZOPANIB HYDROCHLORIDE, PAZOPANIB HYDROCHLORIDE
 PEDIAPRED, PREDNISOLONE SODIUM PHOSPHATE
 PEDMARK, SODIUM THIOSULFATE
 PEG 3350 AND ELECTROLYTES, POLYETHYLENE GLYCOL 3350
 PEG-3350, POTASSIUM CHLORIDE, SODIUM BICARBONATE, SODIUM CHLORIDE, POLYETHYLENE GLYCOL
 PEG-3350, SODIUM SULFATE, SODIUM CHLORIDE, POTASSIUM CHLORIDE, SODIUM ASCORBATE AND
 PEMAZYRE, PEMIGATINIB
 PEMETREXED, PEMETREXED
 PEMETREXED, PEMETREXED DISODIUM
 PEMETREXED DISODIUM, PEMETREXED DISODIUM
 PEMETREXED DITROMETHAMINE, PEMETREXED DITROMETHAMINE
 PEMFEXY, PEMETREXED
 PENCICLOVIR, PENCICLOVIR
 PENICILLAMINE, PENICILLAMINE
 PENICILLIN G POTASSIUM, PENICILLIN G POTASSIUM
 PENICILLIN G POTASSIUM IN PLASTIC CONTAINER, PENICILLIN G POTASSIUM
 PENICILLIN G PROCAINE, PENICILLIN G PROCAINE
 PENICILLIN G SODIUM, PENICILLIN G SODIUM
 PENICILLIN V POTASSIUM, PENICILLIN V POTASSIUM
 PENICILLIN-VK, PENICILLIN V POTASSIUM
 PENTAM, PENTAMIDINE ISETHIONATE
 PENTAMIDINE ISETHIONATE, PENTAMIDINE ISETHIONATE

APPENDIX A - PRODUCT NAME INDEX

** P **

PENTASA, MESALAMINE
PENTOBARBITAL SODIUM, PENTOBARBITAL SODIUM
PENTOLAIR, CYCLOPENTOLATE HYDROCHLORIDE
PENTOSTATIN, PENTOSTATIN
PENTOXIFYLLINE, PENTOXIFYLLINE
PEPCID AC, FAMOTIDINE (OTC)
PEPCID COMPLETE, CALCIUM CARBONATE (OTC)
PERCOCET, ACETAMINOPHEN
PERCODAN, ASPIRIN
PERFOROMIST, FORMOTEROL FUMARATE
PERIDEX, CHLORHEXIDINE GLUCONATE
PERIKABIVEN IN PLASTIC CONTAINER, AMINO ACIDS
PERINDOPRIL ERBUMINE, PERINDOPRIL ERBUMINE
PERIOGARD, CHLORHEXIDINE GLUCONATE
PERMETHRIN, PERMETHRIN (OTC)
PERMETHRIN, PERMETHRIN
PERPHENAZINE, PERPHENAZINE
PERPHENAZINE AND AMITRIPTYLINE HYDROCHLORIDE, AMITRIPTYLINE HYDROCHLORIDE
PERSERIS KIT, RISPERIDONE
PFIZERPEN, PENICILLIN G POTASSIUM
PHEBURANE, SODIUM PHENYLBUTYRATE
PHENDIMETRAZINE TARTRATE, PHENDIMETRAZINE TARTRATE
PHENELZINE SULFATE, PHENELZINE SULFATE
PHENOXYBENZAMINE HYDROCHLORIDE, PHENOXYBENZAMINE HYDROCHLORIDE
PHENTERMINE HYDROCHLORIDE, PHENTERMINE HYDROCHLORIDE
PHENTERMINE HYDROCHLORIDE AND TOPIRAMATE, PHENTERMINE HYDROCHLORIDE
PHENTOLAMINE MESYLATE, PHENTOLAMINE MESYLATE
PHENYLEPHRINE HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE
PHENYLEPHRINE HYDROCHLORIDE AND PROMETHAZINE HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE
PHENYTEK, PHENYTOIN SODIUM
PHENYTOIN, PHENYTOIN
PHENYTOIN SODIUM, PHENYTOIN SODIUM
PHEXXI, CITRIC ACID
PHILITH, ETHINYL ESTRADIOL
PHOSPHOLINE IODIDE, ECHOTHIOPHATE IODIDE
PHOTOFRIN, PORFIMER SODIUM
PHOTREXA, RIBOFLAVIN 5'-PHOSPHATE
PHOTREXA VISCOUS IN DEXTRAN 20%, RIBOFLAVIN 5'-PHOSPHATE
PHOXILLUM B22K 4/0 IN PLASTIC CONTAINER, CALCIUM CHLORIDE
PHOXILLUM BK 4/2.5 IN PLASTIC CONTAINER, CALCIUM CHLORIDE
PHYRAGO, DASATINIB
PHYSIOLYTE IN PLASTIC CONTAINER, MAGNESIUM CHLORIDE
PHYSIOSOL IN PLASTIC CONTAINER, MAGNESIUM CHLORIDE
PHYTONADIONE, PHYTONADIONE
PIFELTRO, DORAVIRINE
PILOCARPINE HYDROCHLORIDE, PILOCARPINE HYDROCHLORIDE
PIMECROLIMUS, PIMECROLIMUS
PIMOZIDE, PIMOZIDE
PIMTREA, DESOGESTREL
PINDOLOL, PINDOLOL
PIOGLITAZONE HYDROCHLORIDE, PIOGLITAZONE HYDROCHLORIDE
PIOGLITAZONE HYDROCHLORIDE AND GLIMEPIRIDE, GLIMEPIRIDE
PIOGLITAZONE HYDROCHLORIDE AND METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
PIPERACILLIN, PIPERACILLIN SODIUM
PIPERACILLIN AND TAZOBACTAM, PIPERACILLIN SODIUM
PIQRAY, ALPELISIB
PIRFENIDONE, PIRFENIDONE
PIROXICAM, PIROXICAM
PITAVASTATIN CALCIUM, PITAVASTATIN CALCIUM
PITOCIN, OXYTOCIN
PIVYA, PIVMECILLINAM HYDROCHLORIDE
PLAN B ONE-STEP, LEVONORGESTREL (OTC)
PLAQUENIL, HYDROXYCHLOROQUINE SULFATE
PLASMA-LYTE 148 IN WATER IN PLASTIC CONTAINER, MAGNESIUM CHLORIDE
PLASMA-LYTE A IN PLASTIC CONTAINER, MAGNESIUM CHLORIDE

APPENDIX A - PRODUCT NAME INDEX

** P **

PLAVIX, CLOPIDOGREL BISULFATE
 PLEGISOL IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 PLENVU, ASCORBIC ACID
 PLERIXAFOR, PLERIXAFOR
 PLUVICTO, LUTETIUM LU-177 VIPIVOTIDE TETRAXETAN
 PODOFILOX, PODOFILOX
 POKONZA, POTASSIUM CHLORIDE
 POLMON, DEXCHLORPHENIRAMINE MALEATE
 POLOCAINE, MEPIVACAINE HYDROCHLORIDE
 POLOCAINE-MPF, MEPIVACAINE HYDROCHLORIDE
 POLYETHYLENE GLYCOL 3350, POLYETHYLENE GLYCOL 3350 (OTC)
 POLYETHYLENE GLYCOL 3350 AND ELECTROLYTES, POLYETHYLENE GLYCOL 3350
 POLYMYXIN B SULFATE, POLYMYXIN B SULFATE
 POMALIDOMIDE, POMALIDOMIDE
 POMALYST, POMALIDOMIDE
 PONSTEL, MEFENAMIC ACID
 PONVORY, PONESIMOD
 PORTIA-28, ETHINYL ESTRADIOL
 POSACONAZOLE, POSACONAZOLE
 POSFREA, PALONOSETRON HYDROCHLORIDE
 POSLUMA, FLOTUFOLASTAT F-18 GALLIUM
 POTASSIUM ACETATE, POTASSIUM ACETATE
 POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
 POTASSIUM CHLORIDE 0.037% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.037% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.037% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.037% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.037% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.037% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.037% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.037% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.075% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.075% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.075% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.075% IN DEXTROSE 3.3% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.075% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.075% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.075% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.075% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.075% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.11% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.11% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.11% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.11% IN DEXTROSE 3.3% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.11% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.11% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.11% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.11% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.11% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.149% IN SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, POTASSIUM
 POTASSIUM CHLORIDE 0.15% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.15% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.15% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.15% IN DEXTROSE 3.3% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.15% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.15% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.15% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.15% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.15% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.15% IN DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
 POTASSIUM CHLORIDE 0.15% IN SODIUM CHLORIDE 0.45%, POTASSIUM CHLORIDE
 POTASSIUM CHLORIDE 0.15% IN SODIUM CHLORIDE 0.9%, POTASSIUM CHLORIDE
 POTASSIUM CHLORIDE 0.15% IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, POTASSIUM CHLORIDE
 POTASSIUM CHLORIDE 0.22% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.22% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER,

APPENDIX A - PRODUCT NAME INDEX

** P **

POTASSIUM CHLORIDE 0.22% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.22% IN DEXTROSE 3.3% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.22% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.22% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.22% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.22% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.22% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.3% AND SODIUM CHLORIDE 0.9%, POTASSIUM CHLORIDE
 POTASSIUM CHLORIDE 0.3% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.3% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.3% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.3% IN DEXTROSE 3.3% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.3% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.3% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.3% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.3% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.3% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.3% IN DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
 POTASSIUM CHLORIDE 10MEQ, POTASSIUM CHLORIDE
 POTASSIUM CHLORIDE 10MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 10MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.225% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 10MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45%, DEXTROSE
 POTASSIUM CHLORIDE 10MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 10MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 10MEQ IN PLASTIC CONTAINER, POTASSIUM CHLORIDE
 POTASSIUM CHLORIDE 15MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 20MEQ, POTASSIUM CHLORIDE
 POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.225% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45%, DEXTROSE
 POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9%, DEXTROSE
 POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
 POTASSIUM CHLORIDE 20MEQ IN PLASTIC CONTAINER, POTASSIUM CHLORIDE
 POTASSIUM CHLORIDE 20MEQ IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, POTASSIUM CHLORIDE
 POTASSIUM CHLORIDE 30MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 30MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45%, DEXTROSE
 POTASSIUM CHLORIDE 30MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 30MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 30MEQ IN PLASTIC CONTAINER, POTASSIUM CHLORIDE
 POTASSIUM CHLORIDE 40MEQ, POTASSIUM CHLORIDE
 POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45%, DEXTROSE
 POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9%, DEXTROSE
 POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 40MEQ IN PLASTIC CONTAINER, POTASSIUM CHLORIDE
 POTASSIUM CHLORIDE 40MEQ IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, POTASSIUM CHLORIDE
 POTASSIUM CHLORIDE 5MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER, CALCIUM
 POTASSIUM CHLORIDE 5MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 5MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE IN PLASTIC CONTAINER, POTASSIUM CHLORIDE
 POTASSIUM CITRATE, POTASSIUM CITRATE
 POTASSIUM IODIDE, POTASSIUM IODIDE (OTC)
 POTASSIUM PHOSPHATES, POTASSIUM PHOSPHATE, DIBASIC
 POTASSIUM PHOSPHATES IN 0.9% SODIUM CHLORIDE, POTASSIUM PHOSPHATE, DIBASIC
 POVIDONE IODINE, POVIDONE-IODINE (OTC)
 PRADAXA, DABIGATRAN ETEXILATE MESYLATE
 PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE
 PRAMOSONE, HYDROCORTISONE ACETATE
 PRASUGREL, PRASUGREL HYDROCHLORIDE
 PRASUGREL HYDROCHLORIDE, PRASUGREL HYDROCHLORIDE
 PRAVASTATIN SODIUM, PRAVASTATIN SODIUM
 PRAZIQUANTEL, PRAZIQUANTEL

APPENDIX A - PRODUCT NAME INDEX

** P **

PRAZOSIN HYDROCHLORIDE, PRAZOSIN HYDROCHLORIDE
PRE-OP, HEXACHLOROPHENE
PRE-OP II, HEXACHLOROPHENE
PRECEDEX, DEXMEDETOMIDINE HYDROCHLORIDE
PRED FORTE, PREDNISOLONE ACETATE
PRED MILD, PREDNISOLONE ACETATE
PREDNICARBATE, PREDNICARBATE
PREDNISOLONE, PREDNISOLONE
PREDNISOLONE ACETATE, PREDNISOLONE ACETATE
PREDNISOLONE SODIUM PHOSPHATE, PREDNISOLONE SODIUM PHOSPHATE
PREDNISON, PREDNISON
PREDNISON INTENSOL, PREDNISON
PREGABALIN, PREGABALIN
PRELONE, PREDNISOLONE
PREMARIN, ESTROGENS, CONJUGATED
PREMASOL 10% IN PLASTIC CONTAINER, AMINO ACIDS
PREMASOL 6% IN PLASTIC CONTAINER, AMINO ACIDS
PREMPHASE 14/14, ESTROGENS, CONJUGATED
PREMPRO, ESTROGENS, CONJUGATED
PREPIDIL, DINOPROSTONE
PRETOMANID, PRETOMANID
PREVACID, LANSOPRAZOLE
PREVACID 24 HR, LANSOPRAZOLE (OTC)
PREVALITE, CHOLESTYRAMINE
PREVANTICS MAXI SWABSTICK, CHLORHEXIDINE GLUCONATE (OTC)
PREVANTICS SWAB, CHLORHEXIDINE GLUCONATE (OTC)
PREVANTICS SWABSTICK, CHLORHEXIDINE GLUCONATE (OTC)
PREVDUO, GLYCOPYRROLATE
PREVIFEM, ETHINYL ESTRADIOL
PREVYMIS, LETERMOVIR
PREZCOBIX, COBICISTAT
PREZISTA, DARUNAVIR
PRIALT, ZICONOTIDE ACETATE
PRIFTIN, RIFAPENTINE
PRILOCAINE HYDROCHLORIDE, PRILOCAINE HYDROCHLORIDE
PRILOCAINE HYDROCHLORIDE AND EPINEPHRINE BITARTRATE, EPINEPHRINE BITARTRATE
PRILOSEC, OMEPRAZOLE MAGNESIUM
PRILOSEC OTC, OMEPRAZOLE MAGNESIUM (OTC)
PRIMAQUINE, PRIMAQUINE PHOSPHATE
PRIMAQUINE PHOSPHATE, PRIMAQUINE PHOSPHATE
PRIMATENE MIST, EPINEPHRINE (OTC)
PRIMAXIN, CILASTATIN SODIUM
PRIMIDONE, PRIMIDONE
PRISMASOL B22GK 4/0 IN PLASTIC CONTAINER, CALCIUM CHLORIDE
PRISMASOL BGK 0/2.5 IN PLASTIC CONTAINER, CALCIUM CHLORIDE
PRISMASOL BGK 2/0 IN PLASTIC CONTAINER, CALCIUM CHLORIDE
PRISMASOL BGK 2/3.5 IN PLASTIC CONTAINER, CALCIUM CHLORIDE
PRISMASOL BGK 4/0/1.2 IN PLASTIC CONTAINER, CALCIUM CHLORIDE
PRISMASOL BGK 4/2.5 IN PLASTIC CONTAINER, CALCIUM CHLORIDE
PRISMASOL BK 0/0/1.2 IN PLASTIC CONTAINER, CALCIUM CHLORIDE
PRISTIQ, DESVENLAFAXINE SUCCINATE
PROAIR HFA, ALBUTEROL SULFATE
PROAIR RESPICLICK, ALBUTEROL SULFATE
PROBALAN, PROBENECID
PROBENECID, PROBENECID
PROBENECID AND COLCHICINE, COLCHICINE
PROCAINAMIDE HYDROCHLORIDE, PROCAINAMIDE HYDROCHLORIDE
PROCARDIA, NIFEDIPINE
PROCARDIA XL, NIFEDIPINE
PROCHLORPERAZINE, PROCHLORPERAZINE
PROCHLORPERAZINE EDISYLATE, PROCHLORPERAZINE EDISYLATE
PROCHLORPERAZINE MALEATE, PROCHLORPERAZINE MALEATE
PROCOMP, PROCHLORPERAZINE MALEATE
PROCTOFOAM HC, HYDROCORTISONE ACETATE
PROCYSBI, CYSTEAMINE BITARTRATE

APPENDIX A - PRODUCT NAME INDEX

** P **

PROGESTERONE, PROGESTERONE
 PROGLYCEM, DIAZOXIDE
 PROGRAF, TACROLIMUS
 PROHANCE, GADOTERIDOL
 PROHANCE MULTIPACK, GADOTERIDOL
 PROLENSA, BROMFENAC SODIUM
 PROMACTA, ELTROMBOPAG OLAMINE
 PROMACTA KIT, ELTROMBOPAG OLAMINE
 PROMETH HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE W/CODEINE PHOSPHATE, CODEINE PHOSPHATE
 PROMETHAZINE DM, DEXTROMETHORPHAN HYDROBROMIDE
 PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
 PROMETHAZINE HYDROCHLORIDE AND CODEINE PHOSPHATE, CODEINE PHOSPHATE
 PROMETHAZINE HYDROCHLORIDE AND DEXTROMETHORPHAN HYDROBROMIDE, DEXTROMETHORPHAN
 PROMETHAZINE HYDROCHLORIDE AND PHENYLEPHRINE HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE
 PROMETHAZINE HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE W/CODEINE PHOSPHATE, CODEINE
 PROMETHAZINE PLAIN, PROMETHAZINE HYDROCHLORIDE
 PROMETHAZINE W/ DEXTROMETHORPHAN, DEXTROMETHORPHAN HYDROBROMIDE
 PROMETHAZINE WITH CODEINE, CODEINE PHOSPHATE
 PROMETHEGAN, PROMETHAZINE HYDROCHLORIDE
 PROMETRIUM, PROGESTERONE
 PROPAFENONE HYDROCHLORIDE, PROPAFENONE HYDROCHLORIDE
 PROPARACAINE HYDROCHLORIDE, PROPARACAINE HYDROCHLORIDE
 PROPECIA, FINASTERIDE
 PROPOFOL, PROPOFOL
 PROPRANOLOL HYDROCHLORIDE, PROPRANOLOL HYDROCHLORIDE
 PROPYLTHIOURACIL, PROPYLTHIOURACIL
 PROSCAR, FINASTERIDE
 PROSOL 20% SULFITE FREE IN PLASTIC CONTAINER, AMINO ACIDS
 PROSTIN VR PEDIATRIC, ALPROSTADIL
 PROTAMINE SULFATE, PROTAMINE SULFATE
 PROTONIX, PANTOPRAZOLE SODIUM
 PROTONIX IV, PANTOPRAZOLE SODIUM
 PROTOPAM CHLORIDE, PRALIDOXIME CHLORIDE
 PROTOPIC, TACROLIMUS
 PROTRIPTYLINE HYDROCHLORIDE, PROTRIPTYLINE HYDROCHLORIDE
 PROVAYBLUE, METHYLENE BLUE
 PROVENTIL-HFA, ALBUTEROL SULFATE
 PROVERA, MEDROXYPROGESTERONE ACETATE
 PROVIGIL, MODAFINIL
 PROVOCHOLINE, METHACHOLINE CHLORIDE
 PROZAC, FLUOXETINE HYDROCHLORIDE
 PRUCALOPRIDE SUCCINATE, PRUCALOPRIDE SUCCINATE
 PSEUDOEPHEDRINE HYDROCHLORIDE, PSEUDOEPHEDRINE HYDROCHLORIDE (OTC)
 PULMICORT FLEXHALER, BUDESONIDE
 PULMICORT RESPULES, BUDESONIDE
 PUR-WASH, PURIFIED WATER (OTC)
 PURIFIED CORTROPHIN GEL, CORTICOTROPIN
 PURINETHOL, MERCAPTOPYRINE
 PURIXAN, MERCAPTOPYRINE
 PYLARIFY, PIFLUFOLASTAT F-18
 PYLERA, BISMUTH SUBCITRATE POTASSIUM
 PYRAZINAMIDE, PYRAZINAMIDE
 PYRIDOSTIGMINE BROMIDE, PYRIDOSTIGMINE BROMIDE
 PYRIDOXINE HYDROCHLORIDE, PYRIDOXINE HYDROCHLORIDE
 PYRIMETHAMINE, PYRIMETHAMINE
 PYRUKYND, MITAPIVAT SULFATE

** Q **

QALSODY, TOFERSEN
 QBRELIS, LISINAPRIL
 QBREXZA, GLYCOPYRRONIUM TOSYLATE
 QELBREE, VILOXAZINE HYDROCHLORIDE
 QINLOCK, RIPRETINIB
 QLOSI, Pilocarpine Hydrochloride
 QNASL, BECLOMETHASONE DIPROPIONATE

APPENDIX A - PRODUCT NAME INDEX

** Q **

QOLIANA, BRIMONIDINE TARTRATE
 QSYMIA, PHENTERMINE HYDROCHLORIDE
 QTERN, DAPAGLIFLOZIN
 QUARTETTE, ETHINYL ESTRADIOL
 QUASENSE, ETHINYL ESTRADIOL
 QUDEXY XR, TOPIRAMATE
 QUELICIN, SUCCINYLCHOLINE CHLORIDE
 QUETIAPINE FUMARATE, QUETIAPINE FUMARATE
 QUILLICHEW ER, METHYLPHENIDATE HYDROCHLORIDE
 QUILLIVANT XR, METHYLPHENIDATE HYDROCHLORIDE
 QUINAPRIL HYDROCHLORIDE, QUINAPRIL HYDROCHLORIDE
 QUINAPRIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 QUINIDINE GLUCONATE, QUINIDINE GLUCONATE
 QUINIDINE SULFATE, QUINIDINE SULFATE
 QUININE SULFATE, QUININE SULFATE
 QULIPTA, ATOGEPANT
 QUTENZA, CAPSAICIN
 QUVIVIQ, DARIDOREXANT HYDROCHLORIDE
 QUZYTTR, CETIRIZINE HYDROCHLORIDE
 QVAR REDIHALER, BECLOMETHASONE DIPROPIONATE

** R **

R-GENE 10, ARGININE HYDROCHLORIDE
 RABEPRAZOLE SODIUM, RABEPRAZOLE SODIUM
 RADICAVA, EDARAVONE
 RADICAVA ORS, EDARAVONE
 RADIOGARDASE (PRUSSIAN BLUE), FERRIC HEXACYANOFERRATE(II)
 RALDESY, TRAZODONE HYDROCHLORIDE
 RALOXIFENE HYDROCHLORIDE, RALOXIFENE HYDROCHLORIDE
 RALTEGRAVIR POTASSIUM, RALTEGRAVIR POTASSIUM
 RAMELTEON, RAMELTEON
 RAMIPRIL, RAMIPRIL
 RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE (OTC)
 RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
 RANOLAZINE, RANOLAZINE
 RAPAFLO, SILODOSIN
 RAPAMUNE, SIROLIMUS
 RAPIBLYK, LANDIOLOL HYDROCHLORIDE
 RAPIVAB, PERAMIVIR
 RASAGILINE MESYLATE, RASAGILINE MESYLATE
 RASUVO, METHOTREXATE
 RAVICTI, GLYCEROL PHENYL BUTYRATE
 RAYALDEE, CALCIFEDIOL
 READI-CAT 2, BARIUM SULFATE
 READI-CAT 2 SMOOTHIE, BARIUM SULFATE
 READYPREP CHG, CHLORHEXIDINE GLUCONATE (OTC)
 RECARBRIO, CILASTATIN SODIUM
 RECLAST, ZOLEDRONIC ACID
 RECORLEV, LEVOKETOCONAZOLE
 RECTIV, NITROGLYCERIN
 REGADENOSON, REGADENOSON
 REGLAN, METOCLOPRAMIDE HYDROCHLORIDE
 REGONOL, PYRIDOSTIGMINE BROMIDE
 RELENZA, ZANAMIVIR
 RELEXXII, METHYLPHENIDATE HYDROCHLORIDE
 RELISTOR, METHYLNALTREXONE BROMIDE
 RELPAX, ELETRIPTAN HYDROBROMIDE
 REMERON, MIRTAZAPINE
 REMERON SOLTAB, MIRTAZAPINE
 REMIFENTANIL HYDROCHLORIDE, REMIFENTANIL HYDROCHLORIDE
 REMODULIN, TREPROSTINIL
 RENACIDIN, CITRIC ACID
 RENAGEL, SEVELAMER HYDROCHLORIDE
 RENOVA, TRETINOIN
 RENVELA, SEVELAMER CARBONATE

APPENDIX A - PRODUCT NAME INDEX

** R **

REPAGLINIDE, REPAGLINIDE
RESTASIS, CYCLOSPORINE
RESTASIS MULTIDOSE, CYCLOSPORINE
RESTORIL, TEMAZEPAM
RETEVMO, SELPERCATINIB
RETIN-A, TRETINOIN
RETIN-A MICRO, TRETINOIN
RETIN-A-MICRO, TRETINOIN
RETISERT, FLUOCINOLONE ACETONIDE
RETROVIR, ZIDOVUDINE
REVATIO, SILDENAFIL CITRATE
REVLIMID, LENALIDOMIDE
REVONTO, DANTROLENE SODIUM
REVUFORJ, REVUMENIB CITRATE
REXTOVY, NALOXONE HYDROCHLORIDE
REXULTI, BREXPIPIRAZOLE
REYATAZ, ATAZANAVIR SULFATE
REYVOW, LASMIDITAN SUCCINATE
REZDIFFRA, RESMETIROM
REZENOPY, NALOXONE HYDROCHLORIDE
REZIPRES, EPHEDRINE HYDROCHLORIDE
REZLIDHIA, OLUTASIDENIB
REZUROCK, BELUMOSUDIL MESYLATE
REZZAYO, REZAFUNGIN ACETATE
RHINOCORT ALLERGY, BUDESONIDE (OTC)
RHOFADÉ, OXYMETAZOLINE HYDROCHLORIDE
RHOPRESSA, NETARSUDIL MESYLATE
RIBAVIRIN, RIBAVIRIN
RIDAURA, AURANOFIN
RIFABUTIN, RIFABUTIN
RIFADIN, RIFAMPIN
RIFAMPIN, RIFAMPIN
RILUZOLE, RILUZOLE
RIMACTANE, RIFAMPIN
RIMANTADINE HYDROCHLORIDE, RIMANTADINE HYDROCHLORIDE
RIMSO-50, DIMETHYL SULFOXIDE
RINGER'S IN PLASTIC CONTAINER, CALCIUM CHLORIDE
RINVOQ, UPADACITINIB
RINVOQ LQ, UPADACITINIB
RIOCIGUAT, RIOCIGUAT
RISEDRONATE SODIUM, RISEDRONATE SODIUM
RISPERDAL, RISPERIDONE
RISPERDAL CONSTA, RISPERIDONE
RISPERIDONE, RISPERIDONE
RITALIN, METHYLPHENIDATE HYDROCHLORIDE
RITALIN LA, METHYLPHENIDATE HYDROCHLORIDE
RITONAVIR, RITONAVIR
RIVASTIGMINE, RIVASTIGMINE
RIVASTIGMINE TARTRATE, RIVASTIGMINE TARTRATE
RIVFLOZA, NEDOSIRAN SODIUM
RIVIVE, NALOXONE HYDROCHLORIDE (OTC)
RIZATRIPTAN BENZOATE, RIZATRIPTAN BENZOATE
ROBAXIN, METHOCARBAMOL
ROBINUL, GLYCOPYRROLATE
ROBINUL FORTE, GLYCOPYRROLATE
ROCALTROL, CALCITRIOL
ROCKLATAN, LATANOPROST
ROCURONIUM BROMIDE, ROCURONIUM BROMIDE
ROFLUMILAST, ROFLUMILAST
ROGAINE (FOR MEN), MINOXIDIL (OTC)
ROGAINE (FOR WOMEN), MINOXIDIL (OTC)
ROGAINE EXTRA STRENGTH (FOR MEN), MINOXIDIL (OTC)
ROMIDEPSIN, ROMIDEPSIN
ROPINIROLE HYDROCHLORIDE, ROPINIROLE HYDROCHLORIDE
ROPIVACAINE HYDROCHLORIDE, ROPIVACAINE HYDROCHLORIDE

APPENDIX A - PRODUCT NAME INDEX

** R **

ROSUVASTATIN CALCIUM, ROSUVASTATIN CALCIUM
 ROWASA, MESALAMINE
 ROXICODONE, OXYCODONE HYDROCHLORIDE
 ROXYBOND, OXYCODONE HYDROCHLORIDE
 ROZEREM, RAMELTEON
 ROZLYTREK, ENTRECTINIB
 RUBRACA, RUCAPARIB CAMSYLATE
 RUBY-FILL, RUBIDIUM CHLORIDE RB-82
 RUFINAMIDE, RUFINAMIDE
 RUKOBIA, FOSTEMSAVIR TROMETHAMINE
 RYALTRIS, MOMETASONE FUROATE
 RYANODEX, DANTROLENE SODIUM
 RYBELSUS, SEMAGLUTIDE
 RYDAPT, MIDOSTAURIN
 RYKINDO, RISPERIDONE
 RYTARY, CARBIDOPA
 RYTELO, IMETELSTAT SODIUM
 RYZUMVI, PHENTOLAMINE MESYLATE

** S **

SABRIL, VIGABATRIN
 SACUBITRIL AND VALSARTAN, SACUBITRIL
 SAFINAMIDE MESYLATE, SAFINAMIDE MESYLATE
 SAFYRAL, DROSPIRENONE
 SALAGEN, PILOCARPINE HYDROCHLORIDE
 SALONPAS, MENTHOL (OTC)
 SAMSCA, TOLVAPTAN
 SANCUSO, GRANISETRON
 SANDIMMUNE, CYCLOSPORINE
 SANDOSTATIN, OCTREOTIDE ACETATE
 SANDOSTATIN LAR, OCTREOTIDE ACETATE
 SAPHRIS, ASENAPINE MALEATE
 SAPROPTERIN DIHYDROCHLORIDE, SAPROPTERIN DIHYDROCHLORIDE
 SAVAYSA, EDOXABAN TOSYLATE
 SAVELLA, MILNACIPRAN HYDROCHLORIDE
 SAXAGLIPTIN, SAXAGLIPTIN HYDROCHLORIDE
 SAXAGLIPTIN AND METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 SAXENDA, LIRAGLUTIDE
 SCANDONEST PLAIN, MEPIVACAINE HYDROCHLORIDE
 SCEMBLIX, ASCIMINIB HYDROCHLORIDE
 SCENESSE, AFAMELANOTIDE
 SCLEROSOL, TALC
 SCOPOLAMINE, SCOPOLAMINE
 SEASONALE, ETHINYL ESTRADIOL
 SEASONIQUE, ETHINYL ESTRADIOL
 SECUADO, ASENAPINE
 SEGLUROMET, ERTUGLIFLOZIN
 SEIZALAM, MIDAZOLAM HYDROCHLORIDE
 SELEGILINE HYDROCHLORIDE, SELEGILINE HYDROCHLORIDE
 SELENIOS ACID, SELENIOS ACID
 SELENIUM SULFIDE, SELENIUM SULFIDE
 SELEXIPAG, SELEXIPAG
 SELZENTRY, MARAVIROC
 SENSORCAINE, BUPIVACAINE HYDROCHLORIDE
 SEPTOCAINE, ARTICAINE HYDROCHLORIDE
 SEPTRA, SULFAMETHOXAZOLE
 SEPTRA DS, SULFAMETHOXAZOLE
 SEREVENT, SALMETEROL XINAFOATE
 SERNIVO, BETAMETHASONE DIPROPIONATE
 SEROMYCIN, CYCLOSERINE
 SEROQUEL, QUETIAPINE FUMARATE
 SEROQUEL XR, QUETIAPINE FUMARATE
 SERTRALINE HYDROCHLORIDE, SERTRALINE HYDROCHLORIDE
 SETLAKIN, ETHINYL ESTRADIOL
 SEVELAMER CARBONATE, SEVELAMER CARBONATE

APPENDIX A - PRODUCT NAME INDEX

** S **

SEVELAMER HYDROCHLORIDE, SEVELAMER HYDROCHLORIDE
SEVOFLURANE, SEVOFLURANE
SEYSARA, SARECYCLINE HYDROCHLORIDE
SEZABY, PHENOBARBITAL SODIUM
SFROWASA, MESALAMINE
SIGNIFOR, PASIREOTIDE DIASPARTATE
SIGNIFOR LAR KIT, PASIREOTIDE PAMOATE
SIKLOS, HYDROXYUREA
SILDENAFIL CITRATE, SILDENAFIL CITRATE
SILENOR, DOXEPIN HYDROCHLORIDE
SILODOSIN, SILODOSIN
SILVADENE, SILVER SULFADIAZINE
SIMBRINZA, BRIMONIDINE TARTRATE
SIMLIYA, DESOGESTREL
SIMPESSE, ETHINYL ESTRADIOL
SIMVASTATIN, SIMVASTATIN
SINCALIDE, SINCALIDE
SINE-AID IB, IBUPROFEN (OTC)
SINEMET, CARBIDOPA
SINGULAIR, MONTELUKAST SODIUM
SINUVA, MOMETASONE FUROATE
SIROLIMUS, SIROLIMUS
SIRTURO, BEDAQUILINE FUMARATE
SITAVIG, ACYCLOVIR
SIVEXTRO, TEDIZOLID PHOSPHATE
SKLICE, IVERMECTIN (OTC)
SKYCLARYS, OMAVELOXOLONE
SKYLA, LEVONORGESTREL
SLYND, DROSPIRENONE
SMOFLIPID 20%, FISH OIL
SOAAZ, TORSEMIDE
SODIUM ACETATE, SODIUM ACETATE
SODIUM BICARBONATE, SODIUM BICARBONATE
SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, SODIUM CHLORIDE
SODIUM CHLORIDE 0.9%, SODIUM CHLORIDE
SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, SODIUM CHLORIDE
SODIUM CHLORIDE 0.9% IN STERILE PLASTIC CONTAINER, SODIUM CHLORIDE
SODIUM CHLORIDE 14.6%, SODIUM CHLORIDE
SODIUM CHLORIDE 23.4%, SODIUM CHLORIDE
SODIUM CHLORIDE 3% IN PLASTIC CONTAINER, SODIUM CHLORIDE
SODIUM CHLORIDE 5% IN PLASTIC CONTAINER, SODIUM CHLORIDE
SODIUM FERRIC GLUCONATE COMPLEX IN SUCROSE, FERRIC OXYHYDROXIDE
SODIUM FLUORIDE F-18, SODIUM FLUORIDE F-18
SODIUM IODIDE I 123, SODIUM IODIDE I-123
SODIUM IODIDE I 131, SODIUM IODIDE I-131
SODIUM NITRITE, SODIUM NITRITE
SODIUM NITROPRUSSIDE, SODIUM NITROPRUSSIDE
SODIUM PHENYLACETATE AND SODIUM BENZOATE, SODIUM BENZOATE
SODIUM PHENYLBUTYRATE, SODIUM PHENYLBUTYRATE
SODIUM PHOSPHATES, SODIUM PHOSPHATE, DIBASIC, HEPTAHYDRATE
SODIUM PHOSPHATES IN PLASTIC CONTAINER, SODIUM PHOSPHATE, DIBASIC, HEPTAHYDRATE
SODIUM PICOSULFATE, MAGNESIUM OXIDE AND ANHYDROUS CITRIC ACID, CITRIC ACID
SODIUM POLYSTYRENE SULFONATE, SODIUM POLYSTYRENE SULFONATE
SODIUM SULFATE, POTASSIUM SULFATE AND MAGNESIUM SULFATE, MAGNESIUM SULFATE
SODIUM TETRADECYL SULFATE, SODIUM TETRADECYL SULFATE
SODIUM THIOSULFATE, SODIUM THIOSULFATE
SOFDRA, SOFPIRONIUM BROMIDE
SOHONOS, PALOVAROTENE
SOJOURN, SEVOFLURANE
SOLIFENACIN SUCCINATE, SOLIFENACIN SUCCINATE
SOLOSEC, SECNIDAZOLE
SOLTAMOX, TAMOXIFEN CITRATE
SOLU-CORTEF, HYDROCORTISONE SODIUM SUCCINATE
SOLU-MEDROL, METHYLPREDNISOLONE SODIUM SUCCINATE
SOLUPREP S, CHLORHEXIDINE GLUCONATE (OTC)

APPENDIX A - PRODUCT NAME INDEX

** S **

SOMA, CARISOPRODOL
SOMATULINE DEPOT, LANREOTIDE ACETATE
SONATA, ZALEPLON
SOOLANTRA, IVERMECTIN
SORAFENIB TOSYLATE, SORAFENIB TOSYLATE
SORBITOL 3% IN PLASTIC CONTAINER, SORBITOL
SORBITOL 3.3% IN PLASTIC CONTAINER, SORBITOL
SORBITOL-MANNITOL IN PLASTIC CONTAINER, MANNITOL
SORILUX, CALCIPOTRIENE
SORINE, SOTALOL HYDROCHLORIDE
SOTALOL HYDROCHLORIDE, SOTALOL HYDROCHLORIDE
SOTRADECOL, SODIUM TETRADECYL SULFATE
SOTYKTU, DEUCRAVACITINIB
SOTYLIZE, SOTALOL HYDROCHLORIDE
SOVALDI, SOFOSBUVIR
SOVUNA, HYDROXYCHLOROQUINE SULFATE
SPINRAZA, NUSINERSEN SODIUM
SPIRIVA, TIOTROPIUM BROMIDE
SPIRIVA RESPIMAT, TIOTROPIUM BROMIDE
SPIRONOLACTONE, SPIRONOLACTONE
SPIRONOLACTONE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
SPORANOX, ITRACONAZOLE
SPRAVATO, ESKETAMINE HYDROCHLORIDE
SPRINTEC, ETHINYL ESTRADIOL
SPRITAM, LEVETIRACETAM
SPRIX, KETOROLAC TROMETHAMINE
SPRYCEL, DASATINIB
SPS, SODIUM POLYSTYRENE SULFONATE
SPY AGENT GREEN KIT, INDOCYANINE GREEN
SSD, SILVER SULFADIAZINE
STALEVO 100, CARBIDOPA
STALEVO 125, CARBIDOPA
STALEVO 150, CARBIDOPA
STALEVO 200, CARBIDOPA
STALEVO 50, CARBIDOPA
STALEVO 75, CARBIDOPA
STEGLATRO, ERTUGLIFLOZIN
STEGLUJAN, ERTUGLIFLOZIN
STENDRA, AVANAFIL
STERILE WATER, STERILE WATER FOR IRRIGATION
STERILE WATER FOR INJECTION, STERILE WATER FOR INJECTION
STERILE WATER FOR INJECTION IN PLASTIC CONTAINER, STERILE WATER FOR INJECTION
STERILE WATER FOR IRRIGATION, STERILE WATER FOR IRRIGATION
STERILE WATER IN PLASTIC CONTAINER, STERILE WATER FOR IRRIGATION
STERITALC, TALC
STIE-CORT, HYDROCORTISONE
STIOLTO RESPIMAT, OLODATEROL HYDROCHLORIDE
STIVARGA, REGORAFENIB
STRATTERA, ATOMOXETINE HYDROCHLORIDE
STREPTOMYCIN SULFATE, STREPTOMYCIN SULFATE
STRIBILD, COBICISTAT
STRIVERDI RESPIMAT, OLODATEROL HYDROCHLORIDE
STROMECTOL, IVERMECTIN
STRONTIUM CHLORIDE SR-89, STRONTIUM CHLORIDE SR-89
SUBLIMAZE PRESERVATIVE FREE, FENTANYL CITRATE
SUBLOCADE, BUPRENORPHINE
SUBOXONE, BUPRENORPHINE HYDROCHLORIDE
SUCCINYLCHOLINE CHLORIDE, SUCCINYLCHOLINE CHLORIDE
SUCRALFATE, SUCRALFATE
SUDAFED 24 HOUR, PSEUDOEPHEDRINE HYDROCHLORIDE (OTC)
SUFENTA PRESERVATIVE FREE, SUFENTANIL CITRATE
SUFENTANIL CITRATE, SUFENTANIL CITRATE
SUFLAVE, MAGNESIUM SULFATE
SUGAMMADEX SODIUM, SUGAMMADEX SODIUM
SULAR, NISOLDIPINE

APPENDIX A - PRODUCT NAME INDEX

** S **

SULFACETAMIDE SODIUM, SULFACETAMIDE SODIUM
 SULFACETAMIDE SODIUM AND PREDNISOLONE SODIUM PHOSPHATE, PREDNISOLONE SODIUM PHOSPHATE
 SULFADIAZINE, SULFADIAZINE
 SULFAMETHOXAZOLE AND TRIMETHOPRIM, SULFAMETHOXAZOLE
 SULFAMYLON, MAFENIDE ACETATE
 SULFASALAZINE, SULFASALAZINE
 SULFATRIM PEDIATRIC, SULFAMETHOXAZOLE
 SULINDAC, SULINDAC
 SUMATRIPTAN, SUMATRIPTAN
 SUMATRIPTAN AND NAPROXEN SODIUM, NAPROXEN SODIUM
 SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
 SUMYCIN, TETRACYCLINE HYDROCHLORIDE
 SUNITINIB MALATE, SUNITINIB MALATE
 SUNLENCA, LENACAPAVIR SODIUM
 SUNOSI, SOLRIAMFETOL HYDROCHLORIDE
 SUPPRELIN LA, HISTRELIN ACETATE
 SUPRANE, DESFLURANE
 SUPRAX, CEFIXIME
 SUPREP BOWEL PREP KIT, MAGNESIUM SULFATE
 SUSTOL, GRANISETRON
 SUTAB, MAGNESIUM SULFATE
 SUTENT, SUNITINIB MALATE
 SYEDA, DROSPIRENONE
 SYFOVRE, PEGCETACOPLAN
 SYMBICORT, BUDESONIDE
 SYMBICORT AEROSPHERE, BUDESONIDE
 SYMBYAX, FLUOXETINE HYDROCHLORIDE
 SYMDEKO (COPACKAGED), IVACAFTOR
 SYMFI, EFAVIRENZ
 SYMFI LO, EFAVIRENZ
 SYMJEPI, EPINEPHRINE
 SYMLIN, PRAMLINTIDE ACETATE
 SYMPAZAN, CLOBAZAM
 SYMPROIC, NALDEMEDINE TOSYLATE
 SYMTUZA, COBICISTAT
 SYNALAR, FLUOCINOLONE ACETONIDE
 SYNAREL, NAFARELIN ACETATE
 SYNJARDY, EMPAGLIFLOZIN
 SYNJARDY XR, EMPAGLIFLOZIN
 SYNTHROID, LEVOTHYROXINE SODIUM **
 SYPRINE, TRIENTINE HYDROCHLORIDE

** T **

TABRECTA, CAPMATINIB HYDROCHLORIDE
 TACLONEX, BETAMETHASONE DIPROPIONATE
 TACROLIMUS, TACROLIMUS
 TADALAFIL, TADALAFIL
 TADLIQ, TADALAFIL
 TAFINLAR, DABRAFENIB MESYLATE
 TAFLUPROST, TAFLUPROST
 TAGAMET HB, CIMETIDINE (OTC)
 TAGITOL V, BARIUM SULFATE
 TAGRISSO, OSIMERTINIB MESYLATE
 TALC, TALC
 TALICIA, AMOXICILLIN
 TALZENNA, TALAZOPARIB TOSYLATE
 TAMIFLU, OSELTAMIVIR PHOSPHATE
 TAMOXIFEN CITRATE, TAMOXIFEN CITRATE
 TAMSULOSIN HYDROCHLORIDE, TAMSULOSIN HYDROCHLORIDE
 TARGRETIN, BEXAROTENE
 TARPEYO, BUDESONIDE
 TASCENSO ODT, FINGOLIMOD LAURYL SULFATE
 TASIGNA, NILOTINIB HYDROCHLORIDE
 TASIMELTEON, TASIMELTEON
 TASMAR, TOLCAPONE

APPENDIX A - PRODUCT NAME INDEX

** T **

TAUVID, FLORTAUCIPIR F-18
TAVABOROLE, TAVABOROLE
TAVALLISSE, FOSTAMATINIB DISODIUM
TAVNEOS, AVACOPAN
TAXOTERE, DOCETAXEL
TAYTULLA, ETHINYL ESTRADIOL
TAZAROTENE, TAZAROTENE
TAZICEF, CEFTAZIDIME
TAZORAC, TAZAROTENE
TAZVERIK, TAZEMETOSTAT HYDROBROMIDE
TECFIDERA, DIMETHYL FUMARATE
TECHNEGAS KIT, TECHNETIUM TC-99M LABELED CARBON
TECHNELITE, TECHNETIUM TC-99M SODIUM PERTECHNETATE GENERATOR
TECHNESCAN, TECHNETIUM TC-99M OXIDRONATE KIT
TECHNESCAN MAG3, TECHNETIUM TC-99M MERTIATIDE KIT
TECHNESCAN PYP KIT, TECHNETIUM TC-99M PYROPHOSPHATE KIT
TECHNETIUM TC 99M MERTIATIDE KIT, TECHNETIUM TC-99M MERTIATIDE KIT
TECHNETIUM TC 99M SESTAMIBI, TECHNETIUM TC-99M SESTAMIBI KIT
TECHNETIUM TC-99M MEBROFENIN, TECHNETIUM TC-99M MEBROFENIN KIT
TECHNETIUM TC-99M MEDRONATE KIT, TECHNETIUM TC-99M MEDRONATE KIT
TECHNETIUM TC-99M SULFUR COLLOID KIT, TECHNETIUM TC-99M SULFUR COLLOID KIT
TECHNETIUM TC99M MERTIATIDE KIT, TECHNETIUM TC-99M MERTIATIDE KIT
TEFLARO, CEFTAROLINE FOSAMIL
TEGRETOL, CARBAMAZEPINE
TEGRETOL-XR, CARBAMAZEPINE
TEKTRUNA, ALISKIREN HEMIFUMARATE
TELMISARTAN, TELMISARTAN
TELMISARTAN AND AMLODIPINE, AMLODIPINE BESYLATE
TELMISARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
TEMAZEPAM, TEMAZEPAM
TEMBEXA, BRINCIDOFIVIR
TEMODAR, TEMOZOLOMIDE
TEMOZOLOMIDE, TEMOZOLOMIDE
TEMSIROLIMUS, TEMSIROLIMUS
TENOFFO VIR ALAFENAMIDE, TENOFFO VIR ALAFENAMIDE FUMARATE
TENOFFO VIR DISOPROXIL FUMARATE, TENOFFO VIR DISOPROXIL FUMARATE
TENORETIC 100, ATENOLOL
TENORETIC 50, ATENOLOL
TENORMIN, ATENOLOL
TEPADINA, THIOTEPA
TEPMETKO, TEPOTINIB HYDROCHLORIDE
TEPYLUTE, THIOTEPA
TERAZOSIN HYDROCHLORIDE, TERAZOSIN HYDROCHLORIDE
TERBINAFINE HYDROCHLORIDE, TERBINAFINE HYDROCHLORIDE (OTC)
TERBINAFINE HYDROCHLORIDE, TERBINAFINE HYDROCHLORIDE
TERBUTALINE SULFATE, TERBUTALINE SULFATE
TERCONAZOLE, TERCONAZOLE
TERIFLUNOMIDE, TERIFLUNOMIDE
TERIL, CARBAMAZEPINE
TERIPARATIDE, TERIPARATIDE
TERLIVAZ, TERLIPRESSIN ACETATE
TESSALON, BENZONATATE
TESTIM, TESTOSTERONE
TESTOPEL, TESTOSTERONE
TESTOSTERONE, TESTOSTERONE
TESTOSTERONE CYPIONATE, TESTOSTERONE CYPIONATE
TESTOSTERONE ENANTHATE, TESTOSTERONE ENANTHATE
TETRABENAZINE, TETRABENAZINE
TETRACAINE HYDROCHLORIDE, TETRACAINE HYDROCHLORIDE
TETRACYCLINE HYDROCHLORIDE, TETRACYCLINE HYDROCHLORIDE
TEXACORT, HYDROCORTISONE
TEZRULY, TERAZOSIN HYDROCHLORIDE
THALITONE, CHLORTHALIDONE
THALLOUS CHLORIDE TL 201, THALLOUS CHLORIDE TL-201
THALOMID, THALIDOMIDE

APPENDIX A - PRODUCT NAME INDEX

** T **

THAM, TROMETHAMINE
THEO-24, THEOPHYLLINE
THEOPHYLLINE, THEOPHYLLINE
THERMAZENE, SILVER SULFADIAZINE
THEROXIDIL, MINOXIDIL (OTC)
THIAMINE HYDROCHLORIDE, THIAMINE HYDROCHLORIDE
THIOGUANINE, THIOGUANINE
THIOLA, TIOPRONIN
THIOLA EC, TIOPRONIN
THIORIDAZINE HYDROCHLORIDE, THIORIDAZINE HYDROCHLORIDE
THIOTEPA, THIOTEPA
THIOTHIXENE, THIOTHIXENE
THYQUIDITY, LEVOTHYROXINE SODIUM
THYRO-TABS, LEVOTHYROXINE SODIUM **
THYROSAFE, POTASSIUM IODIDE (OTC)
TIAGABINE HYDROCHLORIDE, TIAGABINE HYDROCHLORIDE
TIAZAC, DILTIAZEM HYDROCHLORIDE
TIBSOVO, IVOSIDENIB
TICAGRELOR, TICAGRELOR
TICLOPIDINE HYDROCHLORIDE, TICLOPIDINE HYDROCHLORIDE
TIGAN, TRIMETHOBENZAMIDE HYDROCHLORIDE
TIGECYCLINE, TIGECYCLINE
TIGLUTIK KIT, RILUZOLE
TIKOSYN, DOFETILIDE
TIMOLOL, TIMOLOL
TIMOLOL MALEATE, TIMOLOL MALEATE
TIMOPTIC, TIMOLOL MALEATE
TIMOPTIC IN OCUDOSE, TIMOLOL MALEATE
TIMOPTIC-XE, TIMOLOL MALEATE
TINDAMAX, TINIDAZOLE
TINIDAZOLE, TINIDAZOLE
TIOCONAZOLE, TIOCONAZOLE (OTC)
TIOPRONIN, TIOPRONIN
TIOTROPIUM BROMIDE, TIOTROPIUM BROMIDE
TIPIRACIL HYDROCHLORIDE AND TRIFLURIDINE, TIPIRACIL HYDROCHLORIDE
TIROFIBAN HYDROCHLORIDE, TIROFIBAN HYDROCHLORIDE
TIROSINT, LEVOTHYROXINE SODIUM
TIROSINT-SOL, LEVOTHYROXINE SODIUM
TIS-U-SOL, MAGNESIUM SULFATE
TIS-U-SOL IN PLASTIC CONTAINER, MAGNESIUM SULFATE
TISSUEBLUE, BRILLIANT BLUE G
TIVICAY, DOLUTEGRAVIR SODIUM
TIVICAY PD, DOLUTEGRAVIR SODIUM
TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE
TLANDO, TESTOSTERONE UNDECANOATE
TOBI, TOBRAMYCIN
TOBI PODHALER, TOBRAMYCIN
TOBRADEX, DEXAMETHASONE
TOBRADEX ST, DEXAMETHASONE
TOBRAMYCIN, TOBRAMYCIN
TOBRAMYCIN AND DEXAMETHASONE, DEXAMETHASONE
TOBRAMYCIN SULFATE, TOBRAMYCIN SULFATE
TOBRAMYCIN SULFATE (PHARMACY BULK), TOBRAMYCIN SULFATE
TOBREX, TOBRAMYCIN
TOFACITINIB CITRATE, TOFACITINIB CITRATE
TOLAK, FLUOROURACIL
TOLCAPONE, TOLCAPONE
TOLMETIN SODIUM, TOLMETIN SODIUM
TOLSURA, ITRACONAZOLE
TOLTERODINE TARTRATE, TOLTERODINE TARTRATE
TOLVAPTAN, TOLVAPTAN
TOPAMAX, TOPIRAMATE
TOPICORT, DESOXIMETASONE
TOPIRAMATE, TOPIRAMATE
TOPOTECAN HYDROCHLORIDE, TOPOTECAN HYDROCHLORIDE

APPENDIX A - PRODUCT NAME INDEX

** T **

TOPROL-XL, METOPROLOL SUCCINATE
 TOREMIFENE CITRATE, TOREMIFENE CITRATE
 TORISEL, TEMSIROLIMUS
 TORSEMIDE, TORSEMIDE
 TOSYMRA, SUMATRIPTAN
 TOVIAZ, FESOTERODINE FUMARATE
 TPN ELECTROLYTES IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 TPOXX, TECOVIRIMAT
 TRACLEER, BOSENTAN
 TRADJENTA, LINAGLIPTIN
 TRALEMENT, CUPRIC SULFATE
 TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE
 TRAMADOL HYDROCHLORIDE AND ACETAMINOPHEN, ACETAMINOPHEN
 TRANDOLAPRIL, TRANDOLAPRIL
 TRANDOLAPRIL AND VERAPAMIL HYDROCHLORIDE, TRANDOLAPRIL
 TRANEXAMIC ACID, TRANEXAMIC ACID
 TRANSDERM SCOP, SCOPOLAMINE
 TRANXENE, CLORAZEPATE DIPOTASSIUM
 TRANYLCPROMINE SULFATE, TRANYLCPROMINE SULFATE
 TRAVASOL 10% IN PLASTIC CONTAINER, AMINO ACIDS
 TRAVASOL 5.5% IN PLASTIC CONTAINER, AMINO ACIDS
 TRAVASOL 8.5% IN PLASTIC CONTAINER, AMINO ACIDS
 TRAVATAN Z, TRAVOPROST
 TRAVOPROST, TRAVOPROST
 TRAZODONE HYDROCHLORIDE, TRAZODONE HYDROCHLORIDE
 TREANDA, BENDAMUSTINE HYDROCHLORIDE
 TRECATOR, ETHIONAMIDE
 TRELEGY ELLIPTA, FLUTICASONE FUROATE
 TRELSTAR, TRIPTORELIN PAMOATE
 TREPROSTINIL, TREPROSTINIL
 TRETINOIN, TRETINOIN
 TRETINOIN MICROSPHERE, TRETINOIN
 TREXALL, METHOTREXATE SODIUM
 TREXIMET, NAPROXEN SODIUM
 TRI LO SPRINTEC, ETHINYL ESTRADIOL
 TRI-ESTARYLLA, ETHINYL ESTRADIOL
 TRI-LEGEST FE, ETHINYL ESTRADIOL
 TRI-LINYAH, ETHINYL ESTRADIOL
 TRI-LO-ESTARYLLA, ETHINYL ESTRADIOL
 TRI-LO-LINYAH, ETHINYL ESTRADIOL
 TRI-LO-MILI, ETHINYL ESTRADIOL
 TRI-LUMA, FLUOCINOLONE ACETONIDE
 TRI-MILI, ETHINYL ESTRADIOL
 TRI-NORINYL 28-DAY, ETHINYL ESTRADIOL
 TRI-SPRINTEC, ETHINYL ESTRADIOL
 TRIACIN-C, CODEINE PHOSPHATE
 TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE (OTC)
 TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE
 TRIAMTERENE, TRIAMTERENE
 TRIAMTERENE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 TRIAZOLAM, TRIAZOLAM
 TRIBENZOR, AMLODIPINE BESYLATE
 TRIDERM, TRIAMCINOLONE ACETONIDE
 TRIENTINE HYDROCHLORIDE, TRIENTINE HYDROCHLORIDE
 TRISENCE, TRIAMCINOLONE ACETONIDE
 TRIFLUOPERAZINE HYDROCHLORIDE, TRIFLUOPERAZINE HYDROCHLORIDE
 TRIFLURIDINE, TRIFLURIDINE
 TRIHEXYPHENIDYL HYDROCHLORIDE, TRIHEXYPHENIDYL HYDROCHLORIDE
 TRIJARDY XR, EMPAGLIFLOZIN
 TRIKAFTA (COPACKAGED), ELEXACAFTOR, IVACAFTOR, TEZACAFTOR
 TRILEPTAL, OXCARBAZEPINE
 TRILIPIX, CHOLINE FENOFIBRATE
 TRIMETHOBENZAMIDE HYDROCHLORIDE, TRIMETHOBENZAMIDE HYDROCHLORIDE
 TRIMETHOPRIM, TRIMETHOPRIM
 TRIMETHOPRIM SULFATE AND POLYMYXIN B SULFATE, POLYMYXIN B SULFATE

APPENDIX A - PRODUCT NAME INDEX

** T **

TRIMIPRAMINE MALEATE, TRIMIPRAMINE MALEATE
 TRINTELLIX, VORTIOXETINE HYDROBROMIDE
 TRIPTODUR KIT, TRIPTORELIN PAMOATE
 TRISENOX, ARSENIC TRIOXIDE
 TRIUMEQ, ABACAVIR SULFATE
 TRIUMEQ PD, ABACAVIR SULFATE
 TRIVAGIZOLE 3, CLOTRIMAZOLE (OTC)
 TRIVORA-28, ETHINYL ESTRADIOL
 TROKENDI XR, TOPIRAMATE
 TROMETHAMINE, TROMETHAMINE
 TROPHAMINE, AMINO ACIDS
 TROPHAMINE 10%, AMINO ACIDS
 TROPICACYL, TROPICAMIDE
 TROPICAMIDE, TROPICAMIDE
 TROSPIUM CHLORIDE, TROSPIUM CHLORIDE
 TRUDHESA, DIHYDROERGOTAMINE MESYLATE
 TRULANCE, PLECANATIDE
 TRUQAP, CAPIVASERTIB
 TRUVADA, EMTRICITABINE
 TRYNGOLZA (AUTOINJECTOR), OLEZARSEN SODIUM
 TRYVIO, APROCITENTAN
 TUDORZA PRESSAIR, ACLIDINIUM BROMIDE
 TUKYSA, TUCATINIB
 TURALIO, PEXIDARTINIB HYDROCHLORIDE
 TURQOZ, ETHINYL ESTRADIOL
 TUXARIN ER, CHLORPHENIRAMINE MALEATE
 TWIRLA, ETHINYL ESTRADIOL
 TWYNEO, BENZOYL PEROXIDE
 TYBLUME, ETHINYL ESTRADIOL
 TYBOST, COBICISTAT
 TYDEMY, DROSPIRENONE
 TYGACIL, TIGECYCLINE
 TYKERB, LAPATINIB DITOSYLATE
 TYLENOL, ACETAMINOPHEN (OTC)
 TYMLOS, ABALOPARATIDE
 TYRVAYA, VARENICLINE TARTRATE
 TYVASO, TREPROSTINIL
 TYVASO DPI, TREPROSTINIL
 TYZINE, TETRAHYDROZOLINE HYDROCHLORIDE

** U **

UBRELVY, UBROGEPANT
 UCERIS, BUDESONIDE
 ULORIC, FEBUXOSTAT
 ULSPIRA, NITRIC OXIDE
 ULTANE, SEVOFLURANE
 ULTIVA, REMIFENTANIL HYDROCHLORIDE
 ULTRA-TECHNEKOW FM, TECHNETIUM TC-99M SODIUM PERTECHNETATE GENERATOR
 ULTRATAG, TECHNETIUM TC-99M RED BLOOD CELL KIT
 ULTRAVIST (PHARMACY BULK), IOPROMIDE
 ULTRAVIST 300, IOPROMIDE
 ULTRAVIST 370, IOPROMIDE
 UNASYN, AMPICILLIN SODIUM
 UNISOM, DOXYLAMINE SUCCINATE (OTC)
 UNITHROID, LEVOTHYROXINE SODIUM **
 UPNEEQ, OXYMETAZOLINE HYDROCHLORIDE
 UPTRAVI, SELEXIPAG
 UREX, METHENAMINE HIPPURATE
 UROCIT-K, POTASSIUM CITRATE
 UROXATRAL, ALFUZOSIN HYDROCHLORIDE
 URSO 250, URSODIOL
 URSO FORTE, URSODIOL
 URSODIOL, URSODIOL
 UVADEX, METHOXSALLEN
 UZEDY, RISPERIDONE

APPENDIX A - PRODUCT NAME INDEX

** v **

VABOMERE, MEROPENEM
VAFSEO, VADADUSTAT
VAGIFEM, ESTRADIOL
VAGISTAT-1, TIOCONAZOLE (OTC)
VALACYCLOVIR HYDROCHLORIDE, VALACYCLOVIR HYDROCHLORIDE
VALBENAZINE TOSYLATE, VALBENAZINE TOSYLATE
VALCHLOR, MECHLORETHAMINE HYDROCHLORIDE
VALCYTE, VALGANCICLOVIR HYDROCHLORIDE
VALGANCICLOVIR HYDROCHLORIDE, VALGANCICLOVIR HYDROCHLORIDE
VALIUM, DIAZEPAM
VALNAC, BETAMETHASONE VALERATE
VALPROATE SODIUM, VALPROATE SODIUM
VALPROIC ACID, VALPROIC ACID
VALRUBICIN, VALRUBICIN
VALSARTAN, VALSARTAN
VALSARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
VALSTAR PRESERVATIVE FREE, VALRUBICIN
VALTOCO, DIAZEPAM
VALTREX, VALACYCLOVIR HYDROCHLORIDE
VANCOCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE
VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE
VANCOMYCIN HYDROCHLORIDE IN PLASTIC CONTAINER, VANCOMYCIN HYDROCHLORIDE
VANDAZOLE, METRONIDAZOLE
VANFLYTA, QUIZARTINIB DIHYDROCHLORIDE
VANOS, FLUOCINONIDE
VAPRISOL IN 5% DEXTROSE IN PLASTIC CONTAINER, CONIVAPTAN HYDROCHLORIDE
VARDENAFIL HYDROCHLORIDE, VARDENAFIL HYDROCHLORIDE
VARENICLINE TARTRATE, VARENICLINE TARTRATE
VARIBAR HONEY, BARIUM SULFATE
VARIBAR NECTAR, BARIUM SULFATE
VARIBAR PUDDING, BARIUM SULFATE
VARIBAR THIN HONEY, BARIUM SULFATE
VARIBAR THIN LIQUID, BARIUM SULFATE
VARITHENA, POLIDOCANOL
VARUBI, ROLAPITANT HYDROCHLORIDE
VASCEPA, ICOSAPENT ETHYL
VASERETIC, ENALAPRIL MALEATE
VASOPRESSIN, VASOPRESSIN
VASOPRESSIN IN SODIUM CHLORIDE 0.9%, VASOPRESSIN
VASOSTRICT, VASOPRESSIN
VASOTEC, ENALAPRIL MALEATE
VAZALORE, ASPIRIN (OTC)
VAZCULEP, PHENYLEPHRINE HYDROCHLORIDE
VECTICAL, CALCITRIOL
VECURONIUM BROMIDE, VECURONIUM BROMIDE
VEKLURY, REMDESIVIR
VELCADE, BORTEZOMIB
VELETRI, EPOPROSTENOL SODIUM
VELIVET, DESOGESTREL
VELPHORO, FERRIC OXYHYDROXIDE
VELSIPITY, ETRASIMOD ARGININE
VELTASSA, PATIROMER SORBITE X CALCIUM
VELTIN, CLINDAMYCIN PHOSPHATE
VEMLIDY, TENOFOVIR ALAFENAMIDE FUMARATE
VENCLEXTA, VENETOCLAX
VENLAFAXINE BESYLATE, VENLAFAXINE BESYLATE
VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE
VENOFER, FERRIC OXYHYDROXIDE
VENTOLIN HFA, ALBUTEROL SULFATE
VEOZAH, FEZOLINETANT
VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE
VEREGEN, SINECATECHINS
VERELAN, VERAPAMIL HYDROCHLORIDE
VERELAN PM, VERAPAMIL HYDROCHLORIDE
VERKAZIA, CYCLOSPORINE

APPENDIX A - PRODUCT NAME INDEX

** v **

VERQUVO, VERICIGUAT
VERSACLOZ, CLOZAPINE
VERZENIO, ABEMACICLIB
VESICARE, SOLIFENACIN SUCCINATE
VESICARE LS, SOLIFENACIN SUCCINATE
VEVYE, CYCLOSPORINE
VFEND, VORICONAZOLE
VIAGRA, SILDENAFIL CITRATE
VIBATIV, TELAVANCIN HYDROCHLORIDE
VIBERZI, ELUXADOLINE
VIBISONE, CYANOCOBALAMIN
VIBRAMYCIN, DOXYCYCLINE HYCLATE
VICTOZA, LIRAGLUTIDE
VIDAZA, AZACITIDINE
VIENVA, ETHINYL ESTRADIOL
VIGABATRIN, VIGABATRIN
VIGADRONE, VIGABATRIN
VIGAFYDE, VIGABATRIN
VIGAMOX, MOXIFLOXACIN HYDROCHLORIDE
VIGPODER, VIGABATRIN
VIIBRYD, VILAZODONE HYDROCHLORIDE
VIJOICE, ALPELISIB
VILAZODONE HYDROCHLORIDE, VILAZODONE HYDROCHLORIDE
VILTEPSO, VILTOLARSEN
VIMPAT, LACOSAMIDE
VINBLASTINE SULFATE, VINBLASTINE SULFATE
VINCRISTINE SULFATE PFS, VINCRISTINE SULFATE
VINORELBINE TARTRATE, VINORELBINE TARTRATE
VIORELE, DESOGESTREL
VIRACEPT, NELFINAVIR MESYLATE
VIRAZOLE, RIBAVIRIN
VIREAD, TENOFOVIR DISOPROXIL FUMARATE
VIROPTIC, TRIFLURIDINE
VISINE, NAPHAZOLINE HYDROCHLORIDE (OTC)
VISINE L.R., OXYMETAZOLINE HYDROCHLORIDE (OTC)
VISIONBLUE, TRYPAN BLUE
VISIPAQUE 270, IODIXANOL
VISIPAQUE 320, IODIXANOL
VISTARIL, HYDROXYZINE PAMOATE
VISTOGARD, URIDINE TRIACETATE
VISUDYNE, VERTEPORFIN
VITAMIN D, ERGOCALCIFEROL
VITAMIN K1, PHYTONADIONE
VITRAKVI, LAROTRECTINIB SULFATE
VIVACAINE, BUPIVACAINE HYDROCHLORIDE
VIVELLE-DOT, ESTRADIOL
VIVIMUSTA, BENDAMUSTINE HYDROCHLORIDE
VIVITROL, NALTREXONE
VIVJOA, OTESECONAZOLE
VIZAMYL, FLUTEMETAMOL F-18
VIZIMPRO, DACOMITINIB
VOCABRIA, CABOTEGRAVIR SODIUM
VOGELXO, TESTOSTERONE
VOLNEA, DESOGESTREL
VOLTAREN ARTHRITIS PAIN, DICLOFENAC SODIUM (OTC)
VONJO, PACRITINIB CITRATE
VOQUEZNA, VONOPRAZAN FUMARATE
VOQUEZNA DUAL PAK, AMOXICILLIN
VOQUEZNA TRIPLE PAK, AMOXICILLIN
VORANIGO, VORASIDENIB
VORICONAZOLE, VORICONAZOLE
VOSEVI, SOFOSBUVIR
VOSOL HC, ACETIC ACID, GLACIAL
VOTRIENT, PAZOPANIB HYDROCHLORIDE
VOXZOGO, VOSORITIDE

APPENDIX A - PRODUCT NAME INDEX

** V **

VOYDEYA, DANICOPAN
 VRAYLAR, CARIPRAZINE HYDROCHLORIDE
 VTAMA, TAPINAROF
 VUITY, PILOCARPINE HYDROCHLORIDE
 VUMERITY, DIROXIMEL FUMARATE
 VUSION, MICONAZOLE NITRATE
 VYALEV, FOSCARBIDOPA
 VYDUO, NEBIVOLOL HYDROCHLORIDE
 VYFEMLA, ETHINYL ESTRADIOL
 VYLEESI (AUTOINJECTOR), BREMELANOTIDE ACETATE
 VYNDAMAX, TAFAMIDIS
 VYNDAQEL, TAFAMIDIS MEGLUMINE
 VYONDYS 53, GOLODIRSEN
 VYTORIN, EZETIMIBE
 VYVANSE, LISDEXAMFETAMINE DIMESYLATE
 VYXEOS, CYTARABINE
 VYZULTA, LATANOPROSTENE BUNOD

** W **

WAINUA (AUTOINJECTOR), EPLONTERSEN SODIUM
 WAKIX, PITOLISANT HYDROCHLORIDE
 WARFARIN SODIUM, WARFARIN SODIUM
 WEGOVY, SEMAGLUTIDE
 WELCHOL, COLESEVELAM HYDROCHLORIDE
 WELIREG, BELZUTIFAN
 WELLBUTRIN SR, BUPROPION HYDROCHLORIDE
 WELLBUTRIN XL, BUPROPION HYDROCHLORIDE
 WERA, ETHINYL ESTRADIOL
 WINLEVI, CLASCOTERONE
 WIXELA INHUB, FLUTICASONE PROPIONATE
 WOMEN'S ROGAINE, MINOXIDIL (OTC)
 WYNZORA, BETAMETHASONE DIPROPIONATE

** X **

XACDURO (COPACKAGED), DURLOBACTAM SODIUM
 XACIATO, CLINDAMYCIN PHOSPHATE
 XADAGO, SAFINAMIDE MESYLATE
 XALATAN, LATANOPROST
 XALKORI, CRIZOTINIB
 XANAX, ALPRAZOLAM
 XANAX XR, ALPRAZOLAM
 XARELTO, RIVAROXABAN
 XATMEP, METHOTREXATE SODIUM
 XCOPRI, CENOBAMATE
 XDEMVI, LOTILANER
 XELJANZ, TOFACITINIB CITRATE
 XELJANZ XR, TOFACITINIB CITRATE
 XELODA, CAPECITABINE
 XELSTRYM, DEXTROAMPHETAMINE
 XENAZINE, TETRABENAZINE
 XENICAL, ORLISTAT
 XENLETA, LEFAMULIN ACETATE
 XENON XE 133, XENON XE-133
 XENOVIEW, XENON XE-129 HYPERPOLARIZED
 XERAVA, ERAVACYCLINE DIHYDROCHLORIDE
 XERESE, ACYCLOVIR
 XERMELO, TELOTRISTAT ETIPRATE
 XHANCE, FLUTICASONE PROPIONATE
 XIFAXAN, RIFAXIMIN
 XIGDUO XR, DAPAGLIFLOZIN
 XIIDRA, LIFITEGRAST
 XIPERE, TRIAMCINOLONE ACETONIDE
 XOFIGO, RADIUM RA-223 DICHLORIDE
 XOFLUZA, BALOXAVIR MARBOXIL
 XOLREMDI, MAVORIXAFOR

APPENDIX A - PRODUCT NAME INDEX

** X **

XOPENEX HFA, LEVALBUTEROL TARTRATE
 XOSPATA, GILTERITINIB FUMARATE
 XPHOZAH, TENAPANOR HYDROCHLORIDE
 XPOVIO, SELINEXOR
 XROMI, HYDROXYUREA
 XTAMPZA ER, OXYCODONE
 XTANDI, ENZALUTAMIDE
 XULANE, ETHINYL ESTRADIOL
 XURIDEN, URIDINE TRIACETATE
 XYLOCAINE, LIDOCAINE HYDROCHLORIDE
 XYLOCAINE W/ EPINEPHRINE, EPINEPHRINE
 XYOSTED (AUTOINJECTOR), TESTOSTERONE ENANTHATE
 XYREM, SODIUM OXYBATE
 XYWAV, CALCIUM OXYBATE
 XYZAL ALLERGY 24HR, LEVOCETIRIZINE DIHYDROCHLORIDE (OTC)

** Y **

YAELA, DROSPIRENONE
 YARGESA, MIGLUSTAT
 YASMIN, DROSPIRENONE
 YAZ, DROSPIRENONE
 YCANTH, CANTHARIDIN
 YONDELIS, TRABECTEDIN
 YONSA, ABIRATERONE ACETATE
 YORVIPATH, PALOPEGTERIPARATIDE
 YUPELRI, REVEFENACIN
 YUTIQ, FLUOCINOLONE ACETONIDE

** Z **

ZADITOR, KETOTIFEN FUMARATE (OTC)
 ZAFIRLUKAST, ZAFIRLUKAST
 ZALEPLON, ZALEPLON
 ZANAFLEX, TIZANIDINE HYDROCHLORIDE
 ZANOSAR, STREPTOZOCIN
 ZARONTIN, ETHOSUXIMIDE
 ZAVESCA, MIGLUSTAT
 ZAVZPRET, ZAVEGEPANT HYDROCHLORIDE
 ZEGALOGUE, DASIGLUCAGON HYDROCHLORIDE
 ZEGALOGUE (AUTOINJECTOR), DASIGLUCAGON HYDROCHLORIDE
 ZEGERID OTC, OMEPRAZOLE (OTC)
 ZEJULA, NIRAPARIB TOSYLATE
 ZELAPAR, SELEGILINE HYDROCHLORIDE
 ZELBORAF, VEMURAFENIB
 ZELSUVMI, BERDAZIMER SODIUM
 ZEMBRACE SYMTOUCH, SUMATRIPTAN SUCCINATE
 ZEMDRI, PLAZOMICIN SULFATE
 ZEMPLAR, PARICALCITOL
 ZENATANE, ISOTRETINOIN
 ZEPATIER, ELBASVIR
 ZEPBOUND, TIRZEPATIDE
 ZEPBOUND (AUTOINJECTOR), TIRZEPATIDE
 ZEPOSIA, OZANIMOD HYDROCHLORIDE
 ZEPZELCA, LURBINECTEDIN
 ZERBAXA, CEFTOLOZANE SULFATE
 ZERVIAE, CETIRIZINE HYDROCHLORIDE
 ZESTORETIC, HYDROCHLOROTHIAZIDE
 ZESTRIL, LISINAPRIL
 ZETIA, EZETIMIBE
 ZIAC, BISOPROLOL FUMARATE
 ZIAGEN, ABACAVIR SULFATE
 ZIANA, CLINDAMYCIN PHOSPHATE
 ZIDOVUDINE, ZIDOVUDINE
 ZILBRYSQ, ZILUCOPLAN SODIUM
 ZILEUTON, ZILEUTON
 ZILRETTA, TRIAMCINOLONE ACETONIDE

APPENDIX A - PRODUCT NAME INDEX

** Z **

ZILXI, MINOCYCLINE HYDROCHLORIDE
ZIMHI, NALOXONE HYDROCHLORIDE
ZINC CHLORIDE, ZINC CHLORIDE
ZINC CHLORIDE IN PLASTIC CONTAINER, ZINC CHLORIDE
ZINC SULFATE, ZINC SULFATE
ZINGO, LIDOCAINE HYDROCHLORIDE
ZIOPTAN, TAFLUPROST
ZIPRASIDONE HYDROCHLORIDE, ZIPRASIDONE HYDROCHLORIDE
ZIPRASIDONE MESYLATE, ZIPRASIDONE MESYLATE
ZIPSOR, DICLOFENAC POTASSIUM
ZIRGAN, GANCICLOVIR
ZITHROMAX, AZITHROMYCIN
ZITUVIMET, METFORMIN HYDROCHLORIDE
ZITUVIMET XR, METFORMIN HYDROCHLORIDE
ZITUVIO, SITAGLIPTIN
ZOCOR, SIMVASTATIN
ZOKINVY, LONAFARNIB
ZOLADEX, GOSERELIN ACETATE
ZOLEDRONIC, ZOLEDRONIC ACID
ZOLEDRONIC ACID, ZOLEDRONIC ACID
ZOLINZA, VORINOSTAT
ZOLMITRIPTAN, ZOLMITRIPTAN
ZOLOFT, SERTRALINE HYDROCHLORIDE
ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE
ZOMIG, ZOLMITRIPTAN
ZONALON, DOXEPIN HYDROCHLORIDE
ZONEGRAN, ZONISAMIDE
ZONISADE, ZONISAMIDE
ZONISAMIDE, ZONISAMIDE
ZORTRESS, EVEROLIMUS
ZORYVE, ROFLUMILAST
ZOSYN IN PLASTIC CONTAINER, PIPERACILLIN SODIUM
ZOVIA 1/50E-28, ETHINYL ESTRADIOL
ZOVIRAX, ACYCLOVIR
ZTALMY, GANAXOLONE
ZTLIDO, LIDOCAINE
ZUBSOLV, BUPRENORPHINE HYDROCHLORIDE
ZULRESSO, BREXANOLONE
ZUMANDIMINE, DROSPIRENONE
ZUNVEYL, BENZGALANTAMINE GLUCONATE
ZURAGARD, ISOPROPYL ALCOHOL (OTC)
ZURNAI (AUTOINJECTOR), NALMEFENE HYDROCHLORIDE
ZURZUVAE, ZURANOLONE
ZYCLARA, IMIQUIMOD
ZYDELIG, IDELALISIB
ZYFLO, ZILEUTON
ZYKADIA, CERITINIB
ZYLET, LOTEPREDNOL ETABONATE
ZYLOPRIM, ALLOPURINOL
ZYMAXID, GATIFLOXACIN
ZYNRELEF KIT, BUPIVACAINE
ZYPITAMAG, PITAVASTATIN MAGNESIUM
ZYPREXA, OLANZAPINE
ZYPREXA RELPREVV, OLANZAPINE PAMOATE
ZYPREXA ZYDIS, OLANZAPINE
ZYRTEC ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
ZYRTEC-D 12 HOUR, CETIRIZINE HYDROCHLORIDE (OTC)
ZYTIGA, ABIRATERONE ACETATE
ZYVOX, LINEZOLID

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** 3 ******3D IMAGING DRUG**

- * 3D IMAGING DRUG DESIGN AND DEVELOPMENT LLC
AMMONIA N 13, AMMONIA N-13
FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18
SODIUM FLUORIDE F-18, SODIUM FLUORIDE F-18

3M

- * 3M CO
PERIDEX, CHLORHEXIDINE GLUCONATE
- * 3M HEALTH CARE INC
AVAGARD, ALCOHOL (OTC)
DURAPREP, IODINE POVACRYLEX (OTC)

3M HEALTH CARE

- * 3M HEALTH CARE INFECTION PREVENTION DIV
SOLUPREP S, CHLORHEXIDINE GLUCONATE (OTC)

**** 6 ******60 DEGREES PHARMS**

- * 60 DEGREES PHARMACEUTICALS INC
ARAKODA, TAFENOQUINE SUCCINATE

**** A ******AAA USA INC**

- * ADVANCED ACCELERATOR APPLICATIONS USA INC
LUTATHERA, LUTETIUM LU 177 DOTATATE
NETSPOT, GALLIUM DOTATATE GA-68

AADI

- * AADI BIOSCIENCE INC
FYARRO, SIROLIMUS

AAIPHARMA LLC

- * AAIPHARMA LLC
AZASAN, AZATHIOPRINE

ABBVIE

- * ABBVIE
OXYTROL FOR WOMEN, OXYBUTYNIN (OTC)
- * ABBVIE INC
ACULAR LS, KETOROLAC TROMETHAMINE
ACULAR, KETOROLAC TROMETHAMINE
ACUVAIL, KETOROLAC TROMETHAMINE
ACZONE, DAPSONE
ALPHAGAN P, BRIMONIDINE TARTRATE
AVYCAZ, AVIBACTAM SODIUM
CANASA, MESALAMINE
CARAFATE, SUCRALFATE
CELEXA, CITALOPRAM HYDROBROMIDE
COMBIGAN, BRIMONIDINE TARTRATE
CRINONE, PROGESTERONE
CYCLOSPORINE, CYCLOSPORINE
DALVANCE, DALBAVANCIN HYDROCHLORIDE
DEPAKOTE ER, DIVALPROEX SODIUM
DEPAKOTE, DIVALPROEX SODIUM
DUOPA, CARBIDOPA
DURYSTA, BIMATOPROST
FETZIMA, LEVOMILNACIPRAN HYDROCHLORIDE
FML FORTE, FLUOROMETHOLONE
FML, FLUOROMETHOLONE
GENGRAF, CYCLOSPORINE
KALETRA, LOPINAVIR
KYBELLA, DEOXYCHOLIC ACID
LASTACAFT, ALCAFTADINE (OTC)
LATISSE, BIMATOPROST
LEXAPRO, ESCITALOPRAM OXALATE
LINZESS, LINACLOTIDE
LUMIGAN, BIMATOPROST
MAVYRET, GLECAPREVIR
NAMENDA, MEMANTINE HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

* ABBVIE INC
 NAMZARIC, DONEPEZIL HYDROCHLORIDE
 NORVIR, RITONAVIR
 ORIAHNN (COPACKAGED), ELAGOLIX SODIUM, ESTRADIOL, NORETHINDRONE ACETATE
 ORILISSA, ELAGOLIX SODIUM
 OZURDEX, DEXAMETHASONE
 PRED FORTE, PREDNISOLONE ACETATE
 PRED MILD, PREDNISOLONE ACETATE
 QULIPTA, ATOGEPANT
 RAPAFLO, SILODOSIN
 RECTIV, NITROGLYCERIN
 RESTASIS MULTIDOSE, CYCLOSPORINE
 RESTASIS, CYCLOSPORINE
 RINVOQ LQ, UPADACITINIB
 RINVOQ, UPADACITINIB
 SAVELLA, MILNACIPRAN HYDROCHLORIDE
 SYNTHROID, LEVOTHYROXINE SODIUM **
 TEFLARO, CEFTAROLINE FOSAMIL
 TRILIPIX, CHOLINE FENOFIBRATE
 UBRELVY, UBROGEPANT
 ULTANE, SEVOFLURANE
 VENCLEXTA, VENETOCLAX
 VIBERZI, ELUXADOLINE
 VIIBRYD, VILAZODONE HYDROCHLORIDE
 VRAYLAR, CARIPRAZINE HYDROCHLORIDE
 VUITY, PILOCARPINE HYDROCHLORIDE
 VYALEV, FOSCARBIDOPA
 ZEMPLAR, PARICALCITOL
 ZYMAXID, GATIFLOXACIN

ABBVIE ENDOCRINE INC

* ABBVIE ENDOCRINE INC
 LUPRON DEPOT, LEUPROLIDE ACETATE
 LUPRON DEPOT-PED KIT, LEUPROLIDE ACETATE

ABHAI INC

* ABHAI INC
 DEXMETHYLPHENIDATE HYDROCHLORIDE, DEXMETHYLPHENIDATE HYDROCHLORIDE
 METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE

ABHAI LLC

* ABHAI LLC
 ATOVAQUONE, ATOVAQUONE
 BUTALBITAL, ACETAMINOPHEN AND CAFFEINE, ACETAMINOPHEN
 DESMOPRESSIN ACETATE, DESMOPRESSIN ACETATE
 HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
 HYDROCODONE BITARTRATE AND HOMATROPINE METHYLBROMIDE, HOMATROPINE METHYLBROMIDE
 LEFLUNOMIDE, LEFLUNOMIDE
 METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
 OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE
 OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN
 OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
 URSODIOL, URSODIOL

ABON PHARMS LLC

* ABON PHARMACEUTICALS LLC
 ATOVAQUONE, ATOVAQUONE
 GABAPENTIN, GABAPENTIN
 SUCRALFATE, SUCRALFATE

ABRAXIS PHARM

* ABRAXIS PHARMACEUTICAL PRODUCTS
 CLINDAMYCIN PHOSPHATE IN DEXTROSE 5%, CLINDAMYCIN PHOSPHATE

ACACIA

* ACACIA PHARMA LTD
 BARHEMSYS, AMISULPRIDE
 BYFAVO, REMIMAZOLAM BESYLATE

ACADIA PHARMS INC

* ACADIA PHARMACEUTICALS INC
 DAYBUE, TROFINETIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

* ACADIA PHARMACEUTICALS INC
 NUPLAZID, PIMAVANSERIN TARTRATE

ACCELRX LABS

* ACCELRX LABS LLC
 CARISOPRODOL, CARISOPRODOL

ACCORD

* ACCORD BIOPHARMA INC
 CAMCEVI KIT, LEUPROLIDE MESYLATE

ACCORD HLTHCARE

* ACCORD HEALTHCARE INC
 ACETAZOLAMIDE, ACETAZOLAMIDE
 ALLOPURINOL, ALLOPURINOL
 AMITRIPTYLINE HYDROCHLORIDE, AMITRIPTYLINE HYDROCHLORIDE
 AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
 ANASTROZOLE, ANASTROZOLE
 APIXABAN, APIXABAN
 ARGATROBAN IN SODIUM CHLORIDE, ARGATROBAN
 ARIPIPRAZOLE, ARIPIPRAZOLE
 ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM
 ATROPINE SULFATE, ATROPINE SULFATE
 AZACITIDINE, AZACITIDINE
 BENDAMUSTINE HYDROCHLORIDE, BENDAMUSTINE HYDROCHLORIDE
 BICALUTAMIDE, BICALUTAMIDE
 BIVALIRUDIN, BIVALIRUDIN
 BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
 BUSPIRONE HYDROCHLORIDE, BUSPIRONE HYDROCHLORIDE
 CABAZITAXEL, CABAZITAXEL
 CAPECITABINE, CAPECITABINE
 CARBIDOPA AND LEVODOPA, CARBIDOPA
 CARBOPLATIN, CARBOPLATIN
 CINACALCET HYDROCHLORIDE, CINACALCET HYDROCHLORIDE
 CISPLATIN, CISPLATIN
 CLOFARABINE, CLOFARABINE
 CLONAZEPAM, CLONAZEPAM
 CLOZAPINE, CLOZAPINE
 DALFAMPRIDINE, DALFAMPRIDINE
 DAPTOMYCIN, DAPTOMYCIN
 DECITABINE, DECITABINE
 DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE
 DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
 DIMETHYL FUMARATE, DIMETHYL FUMARATE
 DOCETAXEL, DOCETAXEL
 DODEX, CYANOCOBALAMIN
 DOFETILIDE, DOFETILIDE
 DOXAZOSIN MESYLATE, DOXAZOSIN MESYLATE
 EPLERENONE, EPLERENONE
 ESCITALOPRAM OXALATE, ESCITALOPRAM OXALATE
 ETOPOSIDE, ETOPOSIDE
 EZETIMIBE, EZETIMIBE
 FINASTERIDE, FINASTERIDE
 FINGOLIMOD HYDROCHLORIDE, FINGOLIMOD HYDROCHLORIDE
 FLUOROURACIL, FLUOROURACIL
 FULVESTRANT, FULVESTRANT
 FUROSEMIDE, FUROSEMIDE
 GEMCITABINE HYDROCHLORIDE, GEMCITABINE HYDROCHLORIDE
 GLIMEPIRIDE, GLIMEPIRIDE
 GLYCOPYRROLATE, GLYCOPYRROLATE
 HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 HYDROXYCHLOROQUINE SULFATE, HYDROXYCHLOROQUINE SULFATE
 IBANDRONATE SODIUM, IBANDRONATE SODIUM
 IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE
 LETROZOLE, LETROZOLE
 LEVOTHYROXINE SODIUM, LEVOTHYROXINE SODIUM **
 LURASIDONE HYDROCHLORIDE, LURASIDONE HYDROCHLORIDE
 METHOTREXATE SODIUM PRESERVATIVE FREE, METHOTREXATE SODIUM

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

* ACCORD HEALTHCARE INC
 METHOTREXATE SODIUM, METHOTREXATE SODIUM
 METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
 MITOMYCIN, MITOMYCIN
 MONTELUKAST SODIUM, MONTELUKAST SODIUM
 MYCOPHENOLATE MOFETIL, MYCOPHENOLATE MOFETIL
 MYCOPHENOLIC SODIUM, MYCOPHENOLIC SODIUM
 NALOXONE HYDROCHLORIDE, NALOXONE HYDROCHLORIDE
 NALTREXONE HYDROCHLORIDE, NALTREXONE HYDROCHLORIDE
 OLMESARTAN MEDOXOMIL AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 OLMESARTAN MEDOXOMIL, OLMESARTAN MEDOXOMIL
 ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
 OXALIPLATIN, OXALIPLATIN
 OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE
 PACLITAXEL, PACLITAXEL
 PARICALCITOL, PARICALCITOL
 PEMETREXED DISODIUM, PEMETREXED DISODIUM
 PIOGLITAZONE HYDROCHLORIDE, PIOGLITAZONE HYDROCHLORIDE
 PIRFENIDONE, PIRFENIDONE
 PRASUGREL, PRASUGREL HYDROCHLORIDE
 PRAVASTATIN SODIUM, PRAVASTATIN SODIUM
 QUETIAPINE FUMARATE, QUETIAPINE FUMARATE
 REGADENOSON, REGADENOSON
 ROPINIROLE HYDROCHLORIDE, ROPINIROLE HYDROCHLORIDE
 ROSUVASTATIN CALCIUM, ROSUVASTATIN CALCIUM
 SERTRALINE HYDROCHLORIDE, SERTRALINE HYDROCHLORIDE
 SIMVASTATIN, SIMVASTATIN
 SPIRONOLACTONE, SPIRONOLACTONE
 TACROLIMUS, TACROLIMUS
 TADALAFIL, TADALAFIL
 TEMOZOLOMIDE, TEMOZOLOMIDE
 TEMSIROLIMUS, TEMSIROLIMUS
 TERIFLUNOMIDE, TERIFLUNOMIDE
 TOPIRAMATE, TOPIRAMATE
 TOPOTECAN HYDROCHLORIDE, TOPOTECAN HYDROCHLORIDE
 TRAZODONE HYDROCHLORIDE, TRAZODONE HYDROCHLORIDE
 VILAZODONE HYDROCHLORIDE, VILAZODONE HYDROCHLORIDE
 ZOLEDRONIC ACID, ZOLEDRONIC ACID

ACCORD HLTHCARE INC

* ACCORD HEALTHCARE INC USA
 BUSULFAN, BUSULFAN

ACELLA

* ACELLA PHARMACEUTICALS LLC
 AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE
 BETAXOLOL HYDROCHLORIDE, BETAXOLOL HYDROCHLORIDE
 BROMPHENIRAMINE MALEATE, PSEUDOEPHEDRINE HYDROCHLORIDE AND DEXTROMETHORPHAN
 CICLOPIROX, CICLOPIROX
 DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
 DUTASTERIDE, DUTASTERIDE
 LACOSAMIDE, LACOSAMIDE
 NIFEDIPINE, NIFEDIPINE
 PHENYTOIN SODIUM, PHENYTOIN SODIUM

ACELLA PHARMS LLC

* ACELLA PHARMACEUTICALS LLC
 GABAPENTIN, GABAPENTIN

ACER

* ACER THERAPEUTICS INC
 OLPRUVA, SODIUM PHENYLBUTYRATE

ACERUS

* ACERUS PHARMACEUTICALS CORP
 NATESTO, TESTOSTERONE

ACQ PHARMA

* ACQ PHARMA LLC
 MISOPROSTOL, MISOPROSTOL

ACROTECH BIOPHARMA

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

* ACROTECH BIOPHARMA INC
 BELEODAQ, BELINOSTAT
 EVOMELA, MELPHALAN HYDROCHLORIDE
 FOLOTYN, PRALATREXATE
 KHAPZORY, LEVOLEUCOVORIN

ACRUX DDS

* ACRUX DDS PTY LTD
 NITROGLYCERIN, NITROGLYCERIN

ACS DOBFAR

* ACS DOBFAR SPA
 AMPICILLIN AND SULBACTAM, AMPICILLIN SODIUM
 CEFAZOLIN SODIUM, CEFAZOLIN SODIUM
 CEFEPIME HYDROCHLORIDE, CEFEPIME HYDROCHLORIDE
 CEFOXITIN, CEFOXITIN SODIUM
 CEFTAZIDIME, CEFTAZIDIME
 CEFTRIAZONE, CEFTRIAZONE SODIUM
 IMIPENEM AND CILASTATIN, CILASTATIN SODIUM
 MEROPENEM, MEROPENEM

ACS DOBFAR SPA

* ACS DOBFAR SPA
 AMPICILLIN SODIUM, AMPICILLIN SODIUM
 CEFUROXIME SODIUM, CEFUROXIME SODIUM
 ERTAPENEM SODIUM, ERTAPENEM SODIUM
 MEROPENEM, MEROPENEM
 PENICILLIN G POTASSIUM, PENICILLIN G POTASSIUM

ACTAVIS

* ACTAVIS LLC AN INDIRECT WHOLLY OWNED SUB OF TEVA PHARMACEUTICALS USA INC
 DOCETAXEL, DOCETAXEL
 HYDROXOCOBALAMIN, HYDROXOCOBALAMIN
 OXALIPLATIN, OXALIPLATIN
 PEMETREXED, PEMETREXED

ACTAVIS ELIZABETH

* ACTAVIS ELIZABETH LLC
 ALBENDAZOLE, ALBENDAZOLE
 ALPRAZOLAM, ALPRAZOLAM
 BUPRENORPHINE HYDROCHLORIDE AND NALOXONE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE
 BUPRENORPHINE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE
 CARBIDOPA AND LEVODOPA, CARBIDOPA
 CLONAZEPAM, CLONAZEPAM
 CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
 DEFERASIROX, DEFERASIROX
 DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE
 DEXTROAMPHETAMINE SULFATE, DEXTROAMPHETAMINE SULFATE
 DEXTROMETHORPHAN HYDROBROMIDE AND QUINIDINE SULFATE, DEXTROMETHORPHAN HYDROBROMIDE
 DICLOFENAC SODIUM, DICLOFENAC SODIUM
 DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
 DOXEPIN HYDROCHLORIDE, DOXEPIN HYDROCHLORIDE
 DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
 DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE
 FENOFIBRIC ACID, CHOLINE FENOFIBRATE
 FLUVOXAMINE MALEATE, FLUVOXAMINE MALEATE
 GABAPENTIN, GABAPENTIN
 GLYBURIDE AND METFORMIN HYDROCHLORIDE, GLYBURIDE
 GUANFACINE HYDROCHLORIDE, GUANFACINE HYDROCHLORIDE
 HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 INDAPAMIDE, INDAPAMIDE
 LAMOTRIGINE, LAMOTRIGINE
 LISDEXAMFETAMINE DIMESYLATE, LISDEXAMFETAMINE DIMESYLATE
 LOVASTATIN, LOVASTATIN
 METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
 METOPROLOL SUCCINATE, METOPROLOL SUCCINATE
 NIFEDIPINE, NIFEDIPINE
 OXAZEPAM, OXAZEPAM
 OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN
 PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

- * ACTAVIS ELIZABETH LLC
 - PREGABALIN, PREGABALIN
 - PROPRANOLOL HYDROCHLORIDE, PROPRANOLOL HYDROCHLORIDE
 - PROPYLTHIOURACIL, PROPYLTHIOURACIL
 - RANOLAZINE, RANOLAZINE
 - ROPINIROLE HYDROCHLORIDE, ROPINIROLE HYDROCHLORIDE
 - TEMAZEPAM, TEMAZEPAM
- * ACTAVIS ELIZABETH LLC AN INDIRECT WHOLLY OWNED SUB OF TEVA PHARMACEUTICALS USA INC
 - ALPRAZOLAM, ALPRAZOLAM
 - CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
 - MORPHINE SULFATE, MORPHINE SULFATE
 - PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE

ACTAVIS INC

- * ACTAVIS INC
 - DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE
 - DOXORUBICIN HYDROCHLORIDE, DOXORUBICIN HYDROCHLORIDE

ACTAVIS LABS

- * ACTAVIS LABORATORIES INC AN INDIRECT WHOLLY OWNED SUB OF TEVA PHARMACEUTICALS USA INC
 - PERMETHRIN, PERMETHRIN

ACTAVIS LABS FL

- * ACTAVIS LABORATORIES FL INC AN INDIRECT WHOLLY OWNED SUB OF TEVA PHARMACEUTICALS USA INC
 - DESVENLAFAXINE SUCCINATE, DESVENLAFAXINE SUCCINATE
 - GUAIFENESIN AND DEXTROMETHORPHAN HYDROBROMIDE, DEXTROMETHORPHAN HYDROBROMIDE (OTC)
 - GUAIFENESIN AND PSEUDOEPHEDRINE HYDROCHLORIDE, GUAIFENESIN (OTC)
 - GUAIFENESIN, GUAIFENESIN (OTC)
 - ISOTRETINOIN, ISOTRETINOIN
 - MESALAMINE, MESALAMINE
 - METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
 - TOPIRAMATE, TOPIRAMATE

ACTAVIS LABS FL INC

- * ACTAVIS LABORATORIES FL INC
 - BUDESONIDE, BUDESONIDE
 - BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
 - CARTIA XT, DILTIAZEM HYDROCHLORIDE
 - CLARITHROMYCIN, CLARITHROMYCIN
 - CYCLOBENZAPRINE HYDROCHLORIDE, CYCLOBENZAPRINE HYDROCHLORIDE
 - DALFAMPRIDINE, DALFAMPRIDINE
 - DICLOFENAC SODIUM AND MISOPROSTOL, DICLOFENAC SODIUM
 - DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
 - DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
 - DOXYLAMINE SUCCINATE AND PYRIDOXINE HYDROCHLORIDE, DOXYLAMINE SUCCINATE
 - GEFITINIB, GEFITINIB
 - HYDROCODONE BITARTRATE AND IBUPROFEN, HYDROCODONE BITARTRATE
 - LEVETIRACETAM, LEVETIRACETAM
 - METAXALONE, METAXALONE
 - METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 - METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
 - METOPROLOL SUCCINATE, METOPROLOL SUCCINATE
 - NAPROXEN SODIUM, NAPROXEN SODIUM
 - OMEPRAZOLE, OMEPRAZOLE
 - PAROXETINE MESYLATE, PAROXETINE MESYLATE
 - PHENETERMINE HYDROCHLORIDE AND TOPIRAMATE, PHENETERMINE HYDROCHLORIDE
 - POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
 - RAMELTEON, RAMELTEON
 - TAMOXIFEN CITRATE, TAMOXIFEN CITRATE
 - TETRABENAZINE, TETRABENAZINE
 - TRANEXAMIC ACID, TRANEXAMIC ACID
 - TROSPIMUM CHLORIDE, TROSPIMUM CHLORIDE
 - VALGANCICLOVIR HYDROCHLORIDE, VALGANCICLOVIR HYDROCHLORIDE

ACTAVIS LABS UT INC

- * ACTAVIS LABORATORIES UT INC
 - AZELAIC ACID, AZELAIC ACID
 - BUTALBITAL, ACETAMINOPHEN AND CAFFEINE, ACETAMINOPHEN
 - CLINDAMYCIN PHOSPHATE AND BENZOYL PEROXIDE, BENZOYL PEROXIDE
 - CLONIDINE, CLONIDINE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

- * ACTAVIS LABORATORIES UT INC
FIORICET W/ CODEINE, ACETAMINOPHEN
LIDOCAINE, LIDOCAINE
SCOPOLAMINE, SCOPOLAMINE
TESTOSTERONE, TESTOSTERONE
- * ACTAVIS LABORATORIES UT INC INDIRECT WHOLLY OWNED SUB OF TEVA PHARMACEUTICALS USA INC
ADAPALENE AND BENZOYL PEROXIDE, ADAPALENE
PIMECROLIMUS, PIMECROLIMUS
TESTOSTERONE, TESTOSTERONE

ACTAVIS LLC

- * ACTAVIS LLC
AZACITIDINE, AZACITIDINE
DAPSONE, DAPSONE
MELPHALAN HYDROCHLORIDE, MELPHALAN HYDROCHLORIDE
METHYLNALTREXONE BROMIDE, METHYLNALTREXONE BROMIDE

ACTAVIS MID ATLANTIC

- * ACTAVIS MID ATLANTIC LLC
ACYCLOVIR, ACYCLOVIR
ADAPALENE, ADAPALENE
BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
BETAMETHASONE VALERATE, BETAMETHASONE VALERATE
CICLOPIROX, CICLOPIROX
CLINDAMYCIN PHOSPHATE AND TRETINOIN, CLINDAMYCIN PHOSPHATE
CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
CLOTRIMAZOLE AND BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
DESOXIMETASONE, DESOXIMETASONE
DICLOFENAC SODIUM, DICLOFENAC SODIUM
GRISEOFULVIN, GRISEOFULVIN, MICROSIZE
HYDROCODONE BITARTRATE AND HOMATROPINE METHYLBROMIDE, HOMATROPINE METHYLBROMIDE
HYDROCORTISONE BUTYRATE, HYDROCORTISONE BUTYRATE
HYDROCORTISONE, HYDROCORTISONE
LEVETIRACETAM, LEVETIRACETAM
MESALAMINE, MESALAMINE
NITROFURANTOIN, NITROFURANTOIN
NYSTATIN, NYSTATIN
VALNAC, BETAMETHASONE VALERATE
- * ACTAVIS MID ATLANTIC LLC AN INDIRECT WHOLLY OWNED SUB OF TEVA PHARMACEUTICALS USA INC
IBUPROFEN, IBUPROFEN
PERMETHRIN, PERMETHRIN (OTC)

ACTAVIS PHARMA

- * ACTAVIS PHARMA INC AN INDIRECT WHOLLY OWNED SUB OF TEVA PHARMACEUTICALS USA INC
MICONAZOLE NITRATE, MICONAZOLE NITRATE

ACTAVIS TOTOWA

- * ACTAVIS TOTOWA LLC
DESIPRAMINE HYDROCHLORIDE, DESIPRAMINE HYDROCHLORIDE
FLUDARABINE PHOSPHATE, FLUDARABINE PHOSPHATE
IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE
PACLITAXEL, PACLITAXEL
PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
TOPOTECAN HYDROCHLORIDE, TOPOTECAN HYDROCHLORIDE
VINOURELBINE TARTRATE, VINOURELBINE TARTRATE

ACTELION

- * ACTELION PHARMACEUTICALS US INC
OPSUMIT, MACITENTAN
OPSYNVI, MACITENTAN
TRACLEER, BOSENTAN
UPTRAVI, SELEXIPAG
VELETRI, EPOPROSTENOL SODIUM
ZAVESCA, MIGLUSTAT

ADAMIS PHARMS CORP

- * ADAMIS PHARMACEUTICALS CORP
SYMJEPI, EPINEPHRINE

ADAPTIS

- * ADAPTIS PHARMA PRIVATE LTD
ACYCLOVIR, ACYCLOVIR

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

* ADAPTIS PHARMA PRIVATE LTD
 BICALUTAMIDE, BICALUTAMIDE
 DOXEPIN HYDROCHLORIDE, DOXEPIN HYDROCHLORIDE
 DROXIDOPA, DROXIDOPA
 DUTASTERIDE, DUTASTERIDE
 EPLERENONE, EPLERENONE
 ETHACRYNIC ACID, ETHACRYNIC ACID
 ETODOLAC, ETODOLAC
 GLYCOPYRROLATE, GLYCOPYRROLATE
 IBUPROFEN, IBUPROFEN (OTC)
 LURASIDONE HYDROCHLORIDE, LURASIDONE HYDROCHLORIDE
 MOXIFLOXACIN HYDROCHLORIDE, MOXIFLOXACIN HYDROCHLORIDE
 NAPROXEN, NAPROXEN
 NEOSTIGMINE METHYLSULFATE, NEOSTIGMINE METHYLSULFATE
 PREGABALIN, PREGABALIN
 SUCCINYLCHOLINE CHLORIDE, SUCCINYLCHOLINE CHLORIDE
 SUMATRIPTAN, SUMATRIPTAN
 TETRABENAZINE, TETRABENAZINE

ADARE PHARMS INC

* ADARE PHARMACEUTICALS INC
 DEXMETHYLPHENIDATE HYDROCHLORIDE, DEXMETHYLPHENIDATE HYDROCHLORIDE
 POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
 PROPRANOLOL HYDROCHLORIDE, PROPRANOLOL HYDROCHLORIDE

ADIENNE SA

* ADIENNE SA
 TEPADINA, THIOTEPA

ADRASTEIA PHARMA

* ADRASTEIA PHARMA LLC
 COLISTIMETHATE SODIUM, COLISTIMETHATE SODIUM
 HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
 NEOMYCIN SULFATE, NEOMYCIN SULFATE
 NYSTATIN, NYSTATIN
 POLYMYXIN B SULFATE, POLYMYXIN B SULFATE

ADVANZ PHARMA

* ADVANZ PHARMA (US) CORP
 DIBENZYLINE, PHENOXYBENZAMINE HYDROCHLORIDE
 DYRENIUM, TRIAMTERENE
 LANOXIN, DIGOXIN
 NILANDRON, NILUTAMIDE
 ORAPRED ODT, PREDNISOLONE SODIUM PHOSPHATE
 PANRETIN, ALITRETINOIN
 PARNATE, TRANYLCPROMINE SULFATE
 PLAQUENIL, HYDROXYCHLOROQUINE SULFATE
 SALAGEN, PILOCARPINE HYDROCHLORIDE
 UROXATRAL, ALFUZOSIN HYDROCHLORIDE
 ZONEGRAN, ZONISAMIDE

AET PHARMA

* AET PHARMA US INC
 LEFLUNOMIDE, LEFLUNOMIDE
 POSACONAZOLE, POSACONAZOLE

AFAXYS

* AFAXYS PHARMA LLC
 LIDOCAINE HYDROCHLORIDE, LIDOCAINE HYDROCHLORIDE

AFT PHARMS US

* AFT PHARMACEUTICALS US INC
 COMBOGESIC, ACETAMINOPHEN

AGEPHA PHARMA FZ

* AGEPHA PHARMA FZ LLC
 LODOCO, COLCHICINE

AGILE

* AGILE THERAPEUTICS INC
 TWIRLA, ETHINYL ESTRADIOL

AGIOS PHARMS INC

* AGIOS PHARMACEUTICALS INC

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

* AGIOS PHARMACEUTICALS INC
PYRUKYND, MITAPIVAT SULFATE

AGNITIO

* AGNITIO INC
ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM
ETHACRYNIC ACID, ETHACRYNIC ACID
PHYTONADIONE, PHYTONADIONE
TRIAMTERENE, TRIAMTERENE

AGOURON PHARMS

* AGOURON PHARMACEUTICALS LLC
VIRACEPT, NELFINAVIR MESYLATE

AILEX PHARMS LLC

* AILEX PHARMACEUTICALS LLC
BISMUTH SUBSALICYLATE, METRONIDAZOLE AND TETRACYCLINE HYDROCHLORIDE, BISMUTH
CROMOLYN SODIUM, CROMOLYN SODIUM
SODIUM PHENYLACETATE AND SODIUM BENZOATE, SODIUM BENZOATE

AIPING PHARM INC

* AIPING PHARMACEUTICAL INC
AMITRIPTYLINE HYDROCHLORIDE, AMITRIPTYLINE HYDROCHLORIDE
ARFORMOTEROL TARTRATE, ARFORMOTEROL TARTRATE
BENZTROPINE MESYLATE, BENZTROPINE MESYLATE
BUSPIRONE HYDROCHLORIDE, BUSPIRONE HYDROCHLORIDE
CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE
DOXEPIN HYDROCHLORIDE, DOXEPIN HYDROCHLORIDE
ENALAPRIL MALEATE, ENALAPRIL MALEATE
HALOPERIDOL, HALOPERIDOL
LEVETIRACETAM, LEVETIRACETAM
MELOXICAM, MELOXICAM
METOCLOPRAMIDE HYDROCHLORIDE, METOCLOPRAMIDE HYDROCHLORIDE
SORINE, SOTALOL HYDROCHLORIDE

AIRGAS THERAP

* AIRGAS THERAPEUTICS LLC
ULSPIRA, NITRIC OXIDE

AIZANT

* AIZANT DRUG RESEARCH SOLUTIONS PRIVATE LTD
ALBUTEROL SULFATE, ALBUTEROL SULFATE
PIRFENIDONE, PIRFENIDONE
ZILEUTON, ZILEUTON
* AIZANT DRUG RESEARCH SOLUTIONS PVT LTD
ELIGLUSTAT TARTRATE, ELIGLUSTAT TARTRATE

AJANTA PHARMA LTD

* AJANTA PHARMA LTD
ALMOTRIPTAN MALATE, ALMOTRIPTAN MALATE
AMLODIPINE AND OLMESARTAN MEDOXOMIL, AMLODIPINE BESYLATE
ARIPIPRAZOLE, ARIPIPRAZOLE
CAPTOPRIL, CAPTOPRIL
CHLORTHALIDONE, CHLORTHALIDONE
CHOLESTYRAMINE, CHOLESTYRAMINE
CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
DIVALPROEX SODIUM, DIVALPROEX SODIUM
DOXEPIN HYDROCHLORIDE, DOXEPIN HYDROCHLORIDE
DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
DROXIDOPA, DROXIDOPA
DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE
ENTACAPONE, ENTACAPONE
FAMOTIDINE, FAMOTIDINE
FENOFIBRATE (MICRONIZED), FENOFIBRATE
FENOFIBRATE, FENOFIBRATE
FLUPHENAZINE HYDROCHLORIDE, FLUPHENAZINE HYDROCHLORIDE
FLUVOXAMINE MALEATE, FLUVOXAMINE MALEATE
GUANFACINE HYDROCHLORIDE, GUANFACINE HYDROCHLORIDE
LAMOTRIGINE, LAMOTRIGINE
MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
MONTELUKAST SODIUM, MONTELUKAST SODIUM

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ******* AJANTA PHARMA LTD**

NAPROXEN AND ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM
 OMEPRAZOLE AND SODIUM BICARBONATE, OMEPRAZOLE
 OSELTAMIVIR PHOSPHATE, OSELTAMIVIR PHOSPHATE
 OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE
 PALIPERIDONE, PALIPERIDONE
 PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
 PROCHLORPERAZINE MALEATE, PROCHLORPERAZINE MALEATE
 RANOLAZINE, RANOLAZINE
 RISPERIDONE, RISPERIDONE
 SILDENAFIL CITRATE, SILDENAFIL CITRATE
 SILODOSIN, SILODOSIN
 TADALAFIL, TADALAFIL
 TOLTERODINE TARTRATE, TOLTERODINE TARTRATE
 VALGANCICLOVIR HYDROCHLORIDE, VALGANCICLOVIR HYDROCHLORIDE
 VARENICLINE TARTRATE, VARENICLINE TARTRATE
 VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE
 VORICONAZOLE, VORICONAZOLE
 ZOLMITRIPTAN, ZOLMITRIPTAN

AJENAT PHARMS*** AJENAT PHARMACEUTICALS LLC**

DESOXYN, METHAMPHETAMINE HYDROCHLORIDE
 TRANXENE, CLORAZEPATE DIPOTASSIUM

AKARX INC*** AKARX INC**

DOPTELET, AVATROMBOPAG MALEATE

AKEBIA*** AKEBIA THERAPEUTICS INC**

VAFSEO, VADADUSTAT

AKORN*** AKORN OPERATING CO LLC**

FAMOTIDINE, FAMOTIDINE

ALCON*** ALCON LABORATORIES INC**

BSS PLUS, CALCIUM CHLORIDE
 BSS, CALCIUM CHLORIDE
 MIOSTAT, CARBACHOL
 NAPHCN-A, NAPHAZOLINE HYDROCHLORIDE (OTC)

*** ALCON RESEARCH LLC**

PHENYLEPHRINE HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE

ALCON LABS*** ALCON LABORATORIES LTD**

TETRACAINE HYDROCHLORIDE, TETRACAINE HYDROCHLORIDE

ALCON LABS INC*** ALCON LABORATORIES INC**

ALCAINE, PROPARACAINE HYDROCHLORIDE
 CYCLOGYL, CYCLOPENTOLATE HYDROCHLORIDE
 CYCLOMYDRIL, CYCLOPENTOLATE HYDROCHLORIDE
 EYSUVIS, LOTEPRDNOL ETABONATE
 FLUORESCITE, FLUORESCIN SODIUM
 INVELTYS, LOTEPRDNOL ETABONATE
 ISOPTO ATROPINE, ATROPINE SULFATE
 MYDRIACYL, TROPICAMIDE
 PATADAY ONCE DAILY RELIEF, OLOPATADINE HYDROCHLORIDE (OTC)
 PATADAY TWICE DAILY RELIEF, OLOPATADINE HYDROCHLORIDE (OTC)
 RHOPRESSA, NETARSUDIL MESYLATE
 ROCKLATAN, LATANOPROST
 SIMBRINZA, BRIMONIDINE TARTRATE

ALCON PHARMS LTD*** ALCON PHARMACEUTICALS LTD**

BETADINE, POVIDONE-IODINE
 ZADITOR, KETOTIFEN FUMARATE (OTC)

ALEMBIC*** ALEMBIC PHARMACEUTICALS LTD**

ACETAZOLAMIDE, ACETAZOLAMIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

* ALEMBIC PHARMACEUTICALS LTD
 ACITRETIN, ACITRETIN
 ACYCLOVIR, ACYCLOVIR
 ADAPALENE AND BENZOYL PEROXIDE, ADAPALENE
 ADAPALENE, ADAPALENE
 ALBENDAZOLE, ALBENDAZOLE
 ALCAFTADINE, ALCAFTADINE (OTC)
 AMANTADINE HYDROCHLORIDE, AMANTADINE HYDROCHLORIDE
 AMLODIPINE AND OLMESARTAN MEDOXOMIL, AMLODIPINE BESYLATE
 AMLODIPINE BESYLATE AND VALSARTAN, AMLODIPINE BESYLATE
 ARFORMOTEROL TARTRATE, ARFORMOTEROL TARTRATE
 ARIPIPIRAZOLE, ARIPIPIRAZOLE
 ASENAPINE MALEATE, ASENAPINE MALEATE
 AZELASTINE HYDROCHLORIDE, AZELASTINE HYDROCHLORIDE
 AZITHROMYCIN, AZITHROMYCIN
 BEPOTASTINE BESILATE, BEPOTASTINE BESILATE
 BETAMETHASONE VALERATE, BETAMETHASONE VALERATE
 BIMATOPROST, BIMATOPROST
 BISOPROLOL FUMARATE, BISOPROLOL FUMARATE
 BRIMONIDINE TARTRATE AND TIMOLOL MALEATE, BRIMONIDINE TARTRATE
 BRIMONIDINE TARTRATE, BRIMONIDINE TARTRATE
 BROMFENAC SODIUM, BROMFENAC SODIUM
 BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
 CANDESARTAN CILEXETIL, CANDESARTAN CILEXETIL
 CARBIDOPA AND LEVODOPA, CARBIDOPA
 CARBOPROST TROMETHAMINE, CARBOPROST TROMETHAMINE
 CARMUSTINE, CARMUSTINE
 CELECOXIB, CELECOXIB
 CHLORDIAZEPOXIDE HYDROCHLORIDE AND CLIDINIUM BROMIDE, CHLORDIAZEPOXIDE HYDROCHLORIDE
 CHLORPROMAZINE HYDROCHLORIDE, CHLORPROMAZINE HYDROCHLORIDE
 CHLOROTHALIDONE, CHLOROTHALIDONE
 CLARITHROMYCIN, CLARITHROMYCIN
 CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
 CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
 CLOMIPRAMINE HYDROCHLORIDE, CLOMIPRAMINE HYDROCHLORIDE
 CLONAZEPAM, CLONAZEPAM
 CYCLOPHOSPHAMIDE, CYCLOPHOSPHAMIDE
 DABIGATRAN ETEXILATE MESYLATE, DABIGATRAN ETEXILATE MESYLATE
 DAPSONE, DAPSONE
 DARIFENACIN HYDROBROMIDE, DARIFENACIN HYDROBROMIDE
 DEFERASIROX, DEFERASIROX
 DESIPRAMINE HYDROCHLORIDE, DESIPRAMINE HYDROCHLORIDE
 DESONIDE, DESONIDE
 DESVENLAFAXINE SUCCINATE, DESVENLAFAXINE SUCCINATE
 DIAZEPAM, DIAZEPAM
 DICLOFENAC SODIUM, DICLOFENAC SODIUM
 DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
 DIVALPROEX SODIUM, DIVALPROEX SODIUM
 DOCETAXEL, DOCETAXEL
 DOCOSANOL, DOCOSANOL (OTC)
 DORZOLAMIDE HYDROCHLORIDE AND TIMOLOL MALEATE, DORZOLAMIDE HYDROCHLORIDE
 DORZOLAMIDE HYDROCHLORIDE, DORZOLAMIDE HYDROCHLORIDE
 DOXEPIN HYDROCHLORIDE, DOXEPIN HYDROCHLORIDE
 DOXERCALCIFEROL, DOXERCALCIFEROL
 DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
 DOXYCYCLINE, DOXYCYCLINE
 DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE
 ENTACAPONE, ENTACAPONE
 ERLOTINIB HYDROCHLORIDE, ERLOTINIB HYDROCHLORIDE
 ERYTHROMYCIN, ERYTHROMYCIN
 FEBUXOSTAT, FEBUXOSTAT
 FENOFIBRATE (MICRONIZED), FENOFIBRATE
 FENOFIBRATE, FENOFIBRATE
 FENOFIBRIC ACID, CHOLINE FENOFIBRATE
 FESOTERODINE FUMARATE, FESOTERODINE FUMARATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

* ALEMBIC PHARMACEUTICALS LTD
 FLUOROURACIL, FLUOROURACIL
 FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 FLUPHENAZINE HYDROCHLORIDE, FLUPHENAZINE HYDROCHLORIDE
 FORMOTEROL FUMARATE, FORMOTEROL FUMARATE
 FULVESTRANT, FULVESTRANT
 GLYCOPYRROLATE, GLYCOPYRROLATE
 GUANFACINE HYDROCHLORIDE, GUANFACINE HYDROCHLORIDE
 ICATIBANT ACETATE, ICATIBANT ACETATE
 IRBESARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 ITRACONAZOLE, ITRACONAZOLE
 IVABRADINE HYDROCHLORIDE, IVABRADINE HYDROCHLORIDE
 KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
 LACOSAMIDE, LACOSAMIDE
 LAMOTRIGINE, LAMOTRIGINE
 LEVETIRACETAM, LEVETIRACETAM
 LIDOCAINE AND PRILOCAINE, LIDOCAINE
 LIDOCAINE, LIDOCAINE
 LINEZOLID, LINEZOLID
 LURASIDONE HYDROCHLORIDE, LURASIDONE HYDROCHLORIDE
 MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
 MESALAMINE, MESALAMINE
 METHOTREXATE SODIUM, METHOTREXATE SODIUM
 METOLAZONE, METOLAZONE
 METOPROLOL TARTRATE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 METRONIDAZOLE, METRONIDAZOLE
 MIDODRINE HYDROCHLORIDE, MIDODRINE HYDROCHLORIDE
 MODAFINIL, MODAFINIL
 MOXIFLOXACIN HYDROCHLORIDE, MOXIFLOXACIN HYDROCHLORIDE
 MUPIROCIN, MUPIROCIN CALCIUM
 NADOLOL, NADOLOL
 NELARABINE, NELARABINE
 NIFEDIPINE, NIFEDIPINE
 NITROFURANTOIN, NITROFURANTOIN, MACROCRYSTALLINE
 NYSTATIN AND TRIAMCINOLONE ACETONIDE, NYSTATIN
 OLMESARTAN MEDOXOMIL AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 OLMESARTAN MEDOXOMIL, OLMESARTAN MEDOXOMIL
 OLOPATADINE HYDROCHLORIDE, OLOPATADINE HYDROCHLORIDE (OTC)
 OSELTAMIVIR PHOSPHATE, OSELTAMIVIR PHOSPHATE
 PACLITAXEL, PACLITAXEL
 PALIPERIDONE, PALIPERIDONE
 PIRFENIDONE, PIRFENIDONE
 PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE
 PRAZOSIN HYDROCHLORIDE, PRAZOSIN HYDROCHLORIDE
 PREGABALIN, PREGABALIN
 ROPINIROLE HYDROCHLORIDE, ROPINIROLE HYDROCHLORIDE
 SACUBITRIL AND VALSARTAN, SACUBITRIL
 SELEXIPAG, SELEXIPAG
 SOLIFENACIN SUCCINATE, SOLIFENACIN SUCCINATE
 TADALAFIL, TADALAFIL
 TAVABOROLE, TAVABOROLE
 TELMISARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 TELMISARTAN, TELMISARTAN
 TEMAZEPAM, TEMAZEPAM
 TERIFLUNOMIDE, TERIFLUNOMIDE
 THEOPHYLLINE, THEOPHYLLINE
 TIMOLOL MALEATE, TIMOLOL MALEATE
 TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE
 TOBRAMYCIN, TOBRAMYCIN
 TRAVOPROST, TRAVOPROST
 TRETINOIN, TRETINOIN
 VALSARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 VALSARTAN, VALSARTAN
 VARDENAFIL HYDROCHLORIDE, VARDENAFIL HYDROCHLORIDE
 VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

* ALEMBIC PHARMACEUTICALS LTD
 VILAZODONE HYDROCHLORIDE, VILAZODONE HYDROCHLORIDE
 ZOLMITRIPTAN, ZOLMITRIPTAN

ALEMBIC GLOBAL

* ALEMBIC GLOBAL HOLDING SA
 TREPROSTINIL, TREPROSTINIL

ALEMBIC LTD

* ALEMBIC LTD
 LITHIUM CARBONATE, LITHIUM CARBONATE
 ROPINIROLE HYDROCHLORIDE, ROPINIROLE HYDROCHLORIDE

ALEMBIC PHARMS LTD

* ALEMBIC PHARMACEUTICALS LTD
 CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
 DESVENLAFAXINE, DESVENLAFAXINE
 DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
 FAMOTIDINE, FAMOTIDINE
 FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 IRBESARTAN, IRBESARTAN
 LAMOTRIGINE, LAMOTRIGINE
 LEFLUNOMIDE, LEFLUNOMIDE
 LOSARTAN POTASSIUM AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 LOSARTAN POTASSIUM, LOSARTAN POTASSIUM
 MEPROBAMATE, MEPROBAMATE
 METOPROLOL TARTRATE, METOPROLOL TARTRATE
 METRONIDAZOLE, METRONIDAZOLE
 RIVASTIGMINE TARTRATE, RIVASTIGMINE TARTRATE
 VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE

ALEXION PHARMS INC

* ALEXION PHARMACEUTICALS INC
 VOYDEYA, DANICOPAN

ALEXZA PHARMS

* ALEXZA PHARMACEUTICALS INC
 ADASUVE, LOXAPINE

ALIGNSCIENCE PHARMA

* ALIGNSCIENCE PHARMA INC
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 QUETIAPINE FUMARATE, QUETIAPINE FUMARATE

ALIMERA SCIENCES INC

* ALIMERA SCIENCES INC
 ILUVIEN, FLUOCINOLONE ACETONIDE
 YUTIQ, FLUOCINOLONE ACETONIDE

ALKALOIDA ZRT

* ALKALOIDA CHEMICAL CO ZRT
 HYDROXYCHLOROQUINE SULFATE, HYDROXYCHLOROQUINE SULFATE

ALKEM

* ALKEM LABORATORIES LTD
 AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
 GABAPENTIN, GABAPENTIN
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE

ALKEM LABS LTD

* ALKEM LABORATORIES LTD
 AMLODIPINE AND OLMESARTAN MEDOXOMIL, AMLODIPINE BESYLATE
 AMPHETAMINE SULFATE, AMPHETAMINE SULFATE
 APREMILAST, APREMILAST
 ARIPIPRAZOLE, ARIPIPRAZOLE
 ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM
 AZATHIOPRINE, AZATHIOPRINE
 AZELASTINE HYDROCHLORIDE, AZELASTINE HYDROCHLORIDE
 BROMPHENIRAMINE MALEATE, PSEUDOEPHEDRINE HYDROCHLORIDE AND DEXTROMETHORPHAN
 BUPRENORPHINE HYDROCHLORIDE AND NALOXONE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE
 CAPECITABINE, CAPECITABINE
 CEFDINIR, CEFDINIR
 CEFIXIME, CEFIXIME
 CEFPODOXIME PROXETIL, CEFPODOXIME PROXETIL

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

* ALKEM LABORATORIES LTD
 CEFPROZIL, CEFPROZIL
 CEFUROXIME AXETIL, CEFUROXIME AXETIL
 CEPHALEXIN, CEPHALEXIN
 CHLORDIAZEPOXIDE HYDROCHLORIDE AND CLIDINIUM BROMIDE, CHLORDIAZEPOXIDE HYDROCHLORIDE
 CHLORTHALIDONE, CHLORTHALIDONE
 CHOLESTYRAMINE LIGHT, CHOLESTYRAMINE
 CHOLESTYRAMINE, CHOLESTYRAMINE
 CINACALCET HYDROCHLORIDE, CINACALCET HYDROCHLORIDE
 CLOBAZAM, CLOBAZAM
 CLOPIDOGREL BISULFATE, CLOPIDOGREL BISULFATE
 COLCHICINE, COLCHICINE
 COLESEVELAM HYDROCHLORIDE, COLESEVELAM HYDROCHLORIDE
 DABIGATRAN ETEXILATE MESYLATE, DABIGATRAN ETEXILATE MESYLATE
 DALFAMPRIDINE, DALFAMPRIDINE
 DEFERASIROX, DEFERASIROX
 DEXMETHYLPHENIDATE HYDROCHLORIDE, DEXMETHYLPHENIDATE HYDROCHLORIDE
 DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE
 DICLOFENAC POTASSIUM, DICLOFENAC POTASSIUM
 DIMETHYL FUMARATE, DIMETHYL FUMARATE
 DROXIDOPA, DROXIDOPA
 DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE
 ENALAPRIL MALEATE, ENALAPRIL MALEATE
 ERYTHROMYCIN, ERYTHROMYCIN
 ESLICARBAZEPINE ACETATE, ESLICARBAZEPINE ACETATE
 ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM
 EVEROLIMUS, EVEROLIMUS
 EZETIMIBE AND SIMVASTATIN, EZETIMIBE
 EZETIMIBE, EZETIMIBE
 FAMOTIDINE, FAMOTIDINE
 FEBUXOSTAT, FEBUXOSTAT
 FESOTERODINE FUMARATE, FESOTERODINE FUMARATE
 FINASTERIDE, FINASTERIDE
 FINGOLIMOD HYDROCHLORIDE, FINGOLIMOD HYDROCHLORIDE
 FOSFOMYCIN TROMETHAMINE, FOSFOMYCIN TROMETHAMINE
 GABAPENTIN, GABAPENTIN
 HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
 HYDROCHLOROTHIAZIDE AND OLMESARTAN MEDOXOMIL, HYDROCHLOROTHIAZIDE
 IBUPROFEN AND FAMOTIDINE, FAMOTIDINE
 IBUPROFEN, IBUPROFEN
 ITRACONAZOLE, ITRACONAZOLE
 LACOSAMIDE, LACOSAMIDE
 LAMOTRIGINE, LAMOTRIGINE
 LANSOPRAZOLE, LANSOPRAZOLE
 LEVETIRACETAM, LEVETIRACETAM
 LIDOCAINE, LIDOCAINE
 LINEZOLID, LINEZOLID
 LISDEXAMFETAMINE DIMESYLATE, LISDEXAMFETAMINE DIMESYLATE
 LURASIDONE HYDROCHLORIDE, LURASIDONE HYDROCHLORIDE
 MARINOL, DRONABINOL
 MESALAMINE, MESALAMINE
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
 METOPROLOL SUCCINATE, METOPROLOL SUCCINATE
 MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE
 MIRABEGRON, MIRABEGRON
 MORPHINE SULFATE, MORPHINE SULFATE
 MYCOPHENOLATE MOFETIL, MYCOPHENOLATE MOFETIL
 MYCOPHENOLIC SODIUM, MYCOPHENOLIC SODIUM
 NIFEDIPINE, NIFEDIPINE
 OLANZAPINE, OLANZAPINE
 OLMESARTAN MEDOXOMIL, OLMESARTAN MEDOXOMIL
 OXCARBAZEPINE, OXCARBAZEPINE
 OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
 PREGABALIN, PREGABALIN

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ******* ALKEM LABORATORIES LTD**

QUETIAPINE FUMARATE, QUETIAPINE FUMARATE
 RABEPRAZOLE SODIUM, RABEPRAZOLE SODIUM
 RANOLAZINE, RANOLAZINE
 RASAGILINE MESYLATE, RASAGILINE MESYLATE
 RILUZOLE, RILUZOLE
 RIZATRIPTAN BENZOATE, RIZATRIPTAN BENZOATE
 ROFLUMILAST, ROFLUMILAST
 ROSUVASTATIN CALCIUM, ROSUVASTATIN CALCIUM
 RUFINAMIDE, RUFINAMIDE
 SACUBITRIL AND VALSARTAN, SACUBITRIL
 SILDENAFIL CITRATE, SILDENAFIL CITRATE
 SIROLIMUS, SIROLIMUS
 SODIUM SULFATE, POTASSIUM SULFATE AND MAGNESIUM SULFATE, MAGNESIUM SULFATE
 SOLIFENACIN SUCCINATE, SOLIFENACIN SUCCINATE
 TACROLIMUS, TACROLIMUS
 TAMSULOSIN HYDROCHLORIDE, TAMSULOSIN HYDROCHLORIDE
 TEMAZEPAM, TEMAZEPAM
 TICAGRELOR, TICAGRELOR
 TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE
 TOBRAMYCIN, TOBRAMYCIN
 TOLVAPTAN, TOLVAPTAN
 TOPIRAMATE, TOPIRAMATE
 TRAMADOL HYDROCHLORIDE AND ACETAMINOPHEN, ACETAMINOPHEN
 TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE
 VALSARTAN, VALSARTAN
 VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE
 VARENICLINE TARTRATE, VARENICLINE TARTRATE
 VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE
 VIGABATRIN, VIGABATRIN

ALKERMES

* ALKERMES INC
 VIVITROL, NALTREXONE

ALKERMES INC

* ALKERMES INC
 ARISTADA INITIO KIT, ARIPIPRAZOLE LAUROXIL
 ARISTADA, ARIPIPRAZOLE LAUROXIL
 LYBALVI, OLANZAPINE

ALLEGIANCE HLTHCARE

* ALLEGIANCE HEALTHCARE CORP
 POVIDONE IODINE, POVIDONE-IODINE (OTC)

ALLERGAN

* ALLERGAN INC
 AVAGE, TAZAROTENE
 OCUFLOX, OFLOXACIN
 TAZORAC, TAZAROTENE

* ALLERGAN PHARMACEUTICAL
 BETAGAN, LEVOBUNOLOL HYDROCHLORIDE

* ALLERGAN SALES LLC
 BYSTOLIC, NEBIVOLOL HYDROCHLORIDE
 CONDYLOX, PODOFILOX
 ESTRACE, ESTRADIOL
 INFED, FERRIC OXYHYDROXIDE
 OXYTROL, OXYBUTYNIN
 SAPHRIS, ASENAPINE MALEATE
 URSO 250, URSODIOL
 URSO FORTE, URSODIOL

ALLIED

* ALLIED PHARMA INC
 CARISOPRODOL, CARISOPRODOL
 CLOBAZAM, CLOBAZAM
 METHOCARBAMOL, METHOCARBAMOL
 PREDNISONE, PREDNISONE

ALMAJECT

* ALMAJECT INC

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ******* ALMAJECT INC**

CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
 MELPHALAN HYDROCHLORIDE, MELPHALAN HYDROCHLORIDE
 TERIPARATIDE, TERIPARATIDE
 VORICONAZOLE, VORICONAZOLE

ALMATICA*** ALMATICA PHARMA INC**

CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE
 GRALISE, GABAPENTIN
 ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE

*** ALMATICA PHARMA LLC**

LOREEV XR, LORAZEPAM
 MACROBID, NITROFURANTOIN
 MACRODANTIN, NITROFURANTOIN, MACROCRYSTALLINE
 SERTRALINE HYDROCHLORIDE, SERTRALINE HYDROCHLORIDE
 VENLAFAXINE BESYLATE, VENLAFAXINE BESYLATE
 ZESTORETIC, HYDROCHLOROTHIAZIDE

ALMIRALL*** ALMIRALL LLC**

ACZONE, DAPSONE
 AZELEX, AZELAIC ACID
 CORDRAN, FLURANDRENOLIDE
 KLISYRI, TIRBANIBULIN
 SEYSARA, SARECYCLINE HYDROCHLORIDE
 VELTIN, CLINDAMYCIN PHOSPHATE

ALNYLAM PHARMS INC*** ALNYLAM PHARMACEUTICALS INC**

AMVUTTRA, VUTRISIRAN SODIUM
 GIVLAARI, GIVOSIRAN SODIUM
 ONPATTRO, PATISIRAN SODIUM
 OXLUMO, LUMASIRAN SODIUM

ALPHA COGNITION*** ALPHA COGNITION INC**

ZUNVEYL, BENZGALANTAMINE GLUCONATE

ALTAIRE PHARMS INC*** ALTAIRE PHARMACEUTICALS INC**

ALTAFLUOR BENOX, BENOXINATE HYDROCHLORIDE
 CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE
 DICLOFENAC SODIUM, DICLOFENAC SODIUM
 NAPHAZOLINE HYDROCHLORIDE AND PHENIRAMINE MALEATE, NAPHAZOLINE HYDROCHLORIDE (OTC)
 OFLOXACIN, OFLOXACIN

ALTATHERA PHARMS LLC*** ALTATHERA PHARMACEUTICALS LLC**

SOTALOL HYDROCHLORIDE, SOTALOL HYDROCHLORIDE

ALTHERA PHARMS*** ALTHERA PHARMACEUTICALS LLC**

LYPQOZET, ATORVASTATIN CALCIUM

ALVOGEN*** ALVOGEN INC**

ACETYLCYSTEINE, ACETYLCYSTEINE
 BONSIITY, TERIPARATIDE
 BUPRENORPHINE HYDROCHLORIDE AND NALOXONE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE
 BUPRENORPHINE, BUPRENORPHINE
 CARBIDOPA, CARBIDOPA
 DEXAMETHASONE, DEXAMETHASONE
 DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE
 DISULFIRAM, DISULFIRAM
 ESTRADIOL, ESTRADIOL
 FELBAMATE, FELBAMATE
 HYDROCODONE BITARTRATE, HYDROCODONE BITARTRATE
 OSELTAMIVIR PHOSPHATE, OSELTAMIVIR PHOSPHATE
 OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN
 OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
 PREGABALIN, PREGABALIN
 PYRIDOSTIGMINE BROMIDE, PYRIDOSTIGMINE BROMIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ******* ALVOGEN INC**

PYRIMETHAMINE, PYRIMETHAMINE
 RIVASTIGMINE, RIVASTIGMINE
 THYRO-TABS, LEVOTHYROXINE SODIUM **
 UREX, METHENAMINE HIPPURATE

AM REAGENT*** AMERICAN REAGENT INC**

ACETYLCYSTEINE, ACETYLCYSTEINE
 ATROPINE SULFATE, ATROPINE SULFATE
 BETAMETHASONE ACETATE AND BETAMETHASONE SODIUM PHOSPHATE, BETAMETHASONE ACETATE
 CAFFEINE CITRATE, CAFFEINE CITRATE
 CALCIUM CHLORIDE 10%, CALCIUM CHLORIDE
 CHLOROTHIAZIDE SODIUM, CHLOROTHIAZIDE SODIUM
 CUPRIC SULFATE, CUPRIC SULFATE
 CYANOCOBALAMIN, CYANOCOBALAMIN
 DICYCLOMINE HYDROCHLORIDE, DICYCLOMINE HYDROCHLORIDE
 DROPERIDOL, DROPERIDOL
 EPINEPHRINE, EPINEPHRINE
 ESTRADIOL VALERATE, ESTRADIOL VALERATE
 FOMEPIZOLE, FOMEPIZOLE
 GLYCOPYRROLATE, GLYCOPYRROLATE
 HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
 HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE
 INJECTAFER, FERRIC CARBOXYMALTOSIDE
 ISOSULFAN BLUE, ISOSULFAN BLUE
 LEVOCARNITINE, LEVOCARNITINE
 METHOCARBAMOL, METHOCARBAMOL
 METHYLERGONOVINE MALEATE, METHYLERGONOVINE MALEATE
 MULTRYS, CUPRIC SULFATE
 NICARDIPINE HYDROCHLORIDE, NICARDIPINE HYDROCHLORIDE
 NITROGLYCERIN, NITROGLYCERIN
 OLANZAPINE, OLANZAPINE
 PACLITAXEL, PACLITAXEL
 POTASSIUM PHOSPHATES, POTASSIUM PHOSPHATE, DIBASIC
 SELENIOS ACID, SELENIOS ACID
 SODIUM PHOSPHATES, SODIUM PHOSPHATE, DIBASIC, HEPTAHYDRATE
 STERILE WATER FOR INJECTION, STERILE WATER FOR INJECTION
 TESTOSTERONE CYPIONATE, TESTOSTERONE CYPIONATE
 TRALEMENT, CUPRIC SULFATE
 TRANEXAMIC ACID, TRANEXAMIC ACID
 VASOPRESSIN, VASOPRESSIN
 VENOFER, FERRIC OXYHYDROXIDE
 ZINC SULFATE, ZINC SULFATE

AMARIN PHARMS*** AMARIN PHARMACEUTICALS IRELAND LTD**

VASCEPA, ICOSAPENT ETHYL

AMGEN INC*** AMGEN INC**

CORLANOR, IVABRADINE
 CORLANOR, IVABRADINE HYDROCHLORIDE
 LUMAKRAS, SOTORASIB
 OTEZLA, APREMILAST

AMICI PHARMA*** AMICI PHARMA INC**

CATAFLAM, DICLOFENAC POTASSIUM
 DIGOXIN, DIGOXIN

AMICUS THERAP US*** AMICUS THERAPEUTICS US LLC**

GALAFOLD, MIGALASTAT HYDROCHLORIDE
 OPFOLDA, MIGLUSTAT

AMIVAS*** AMIVAS INC**

ARTESUNATE, ARTESUNATE

AMNEAL*** AMNEAL EU LTD**

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ******* AMNEAL EU LTD**

AMINOCAPROIC ACID, AMINOCAPROIC ACID
 ARSENIC TRIOXIDE, ARSENIC TRIOXIDE
 ATROPINE SULFATE, ATROPINE SULFATE
 AZACITIDINE, AZACITIDINE
 BRIMONIDINE TARTRATE AND TIMOLOL MALEATE, BRIMONIDINE TARTRATE
 BROMFENAC SODIUM, BROMFENAC SODIUM
 BUSULFAN, BUSULFAN
 CALCIUM CHLORIDE 10%, CALCIUM CHLORIDE
 CALCIUM GLUCONATE IN SODIUM CHLORIDE, CALCIUM GLUCONATE
 CALCIUM GLUCONATE, CALCIUM GLUCONATE
 CARBOPROST TROMETHAMINE, CARBOPROST TROMETHAMINE
 CARMUSTINE, CARMUSTINE
 CHLORDIAZEPOXIDE HYDROCHLORIDE AND CLIDINIUM BROMIDE, CHLORDIAZEPOXIDE HYDROCHLORIDE
 CIPROFLOXACIN; DEXAMETHASONE, CIPROFLOXACIN
 CLOFARABINE, CLOFARABINE
 CYCLOPHOSPHAMIDE, CYCLOPHOSPHAMIDE
 DARUNAVIR, DARUNAVIR
 DEXAMETHASONE SODIUM PHOSPHATE PRESERVATIVE FREE, DEXAMETHASONE SODIUM PHOSPHATE
 DEXAMETHASONE SODIUM PHOSPHATE, DEXAMETHASONE SODIUM PHOSPHATE
 DEXAMETHASONE, DEXAMETHASONE
 DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE
 DIFLUPREDNATE, DIFLUPREDNATE
 DIMETHYL FUMARATE, DIMETHYL FUMARATE
 DOCETAXEL, DOCETAXEL
 DOXORUBICIN HYDROCHLORIDE, DOXORUBICIN HYDROCHLORIDE
 DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
 EPHEDRINE SULFATE, EPHEDRINE SULFATE
 ESMOLOL HYDROCHLORIDE, ESMOLOL HYDROCHLORIDE
 ETRAVIRINE, ETRAVIRINE
 EXENATIDE SYNTHETIC, EXENATIDE SYNTHETIC
 FLUOROMETHOLONE, FLUOROMETHOLONE
 FLUPHENAZINE HYDROCHLORIDE, FLUPHENAZINE HYDROCHLORIDE
 FOSCARNET SODIUM, FOSCARNET SODIUM
 FOSFOMYCIN TROMETHAMINE, FOSFOMYCIN TROMETHAMINE
 FULVESTRANT, FULVESTRANT
 GLYCOPYRROLATE, GLYCOPYRROLATE
 GRANISETRON HYDROCHLORIDE, GRANISETRON HYDROCHLORIDE
 ISOPROTERENOL HYDROCHLORIDE, ISOPROTERENOL HYDROCHLORIDE
 LEUPROLIDE ACETATE, LEUPROLIDE ACETATE
 LEVOLEUCOVORIN CALCIUM, LEVOLEUCOVORIN CALCIUM
 LOTE Prednol Etabonate, LOTE Prednol Etabonate
 MAGNESIUM SULFATE IN PLASTIC CONTAINER, MAGNESIUM SULFATE
 MEDROXYPROGESTERONE ACETATE, MEDROXYPROGESTERONE ACETATE
 METHYLPREDNISOLONE ACETATE, METHYLPREDNISOLONE ACETATE
 METHYLPREDNISOLONE SODIUM SUCCINATE, METHYLPREDNISOLONE SODIUM SUCCINATE
 METRONIDAZOLE IN PLASTIC CONTAINER, METRONIDAZOLE
 METYROSINE, METYROSINE
 NELARABINE, NELARABINE
 NEOSTIGMINE METHYLSULFATE, NEOSTIGMINE METHYLSULFATE
 NICARDIPINE HYDROCHLORIDE, NICARDIPINE HYDROCHLORIDE
 NOREPINEPHRINE BITARTRATE, NOREPINEPHRINE BITARTRATE
 OFLOXACIN, OFLOXACIN
 PHENOXYBENZAMINE HYDROCHLORIDE, PHENOXYBENZAMINE HYDROCHLORIDE
 PHENYLEPHRINE HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE
 PILOCARPINE HYDROCHLORIDE, PILOCARPINE HYDROCHLORIDE
 PIRFENIDONE, PIRFENIDONE
 PLERIXAFOR, PLERIXAFOR
 POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
 POTASSIUM PHOSPHATES IN 0.9% SODIUM CHLORIDE, POTASSIUM PHOSPHATE, DIBASIC
 POTASSIUM PHOSPHATES, POTASSIUM PHOSPHATE, DIBASIC
 PREDNISONE, PREDNISONE
 PROCHLORPERAZINE EDISYLATE, PROCHLORPERAZINE EDISYLATE
 PROCHLORPERAZINE MALEATE, PROCHLORPERAZINE MALEATE
 PROPOFOL, PROPOFOL

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

* AMNEAL EU LTD
 ROPIVACAINE HYDROCHLORIDE, ROPIVACAINE HYDROCHLORIDE
 SODIUM NITROPRUSSIDE, SODIUM NITROPRUSSIDE
 SUCCINYLCHOLINE CHLORIDE, SUCCINYLCHOLINE CHLORIDE
 THEOPHYLLINE, THEOPHYLLINE
 THIOTHIXENE, THIOTHIXENE
 TIOPRONIN, TIOPRONIN
 TRANEXAMIC ACID, TRANEXAMIC ACID
 TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE
 VASOPRESSIN, VASOPRESSIN
 VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE
 ZOLEDRONIC ACID, ZOLEDRONIC ACID

* AMNEAL PHARMACEUTICALS LLC
 ACTIVELLA, ESTRADIOL
 ACYCLOVIR, ACYCLOVIR
 AMINOCAPROIC ACID, AMINOCAPROIC ACID
 AZATHIOPRINE, AZATHIOPRINE
 AZELASTINE HYDROCHLORIDE, AZELASTINE HYDROCHLORIDE
 AZELASTINE HYDROCHLORIDE, AZELASTINE HYDROCHLORIDE (OTC)
 AZITHROMYCIN, AZITHROMYCIN
 BACLOFEN, BACLOFEN
 BEXAROTENE, BEXAROTENE
 BUPRENORPHINE, BUPRENORPHINE
 CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE
 CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
 CLOBAZAM, CLOBAZAM
 CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
 DAPSONE, DAPSONE
 DEXTROMETHORPHAN POLISTIREX, DEXTROMETHORPHAN POLISTIREX (OTC)
 DICLOFENAC SODIUM, DICLOFENAC SODIUM
 DOXEPIN HYDROCHLORIDE, DOXEPIN HYDROCHLORIDE
 ELURYNG, ETHINYL ESTRADIOL
 ENALAPRIL MALEATE, ENALAPRIL MALEATE
 ESTRADIOL, ESTRADIOL
 ETHINYL ESTRADIOL AND NORELGESTROMIN, ETHINYL ESTRADIOL
 FENOFIBRATE, FENOFIBRATE
 FLUOCINONIDE, FLUOCINONIDE
 IPRATROPIUM BROMIDE, IPRATROPIUM BROMIDE
 LIDOCAINE, LIDOCAINE
 LIORESAL, BACLOFEN
 LISDEXAMFETAMINE DIMESYLATE, LISDEXAMFETAMINE DIMESYLATE
 LUBIPROSTONE, LUBIPROSTONE
 LYVISPAH, BACLOFEN
 MUPIROCIN, MUPIROCIN CALCIUM
 MYCOPHENOLATE MOFETIL, MYCOPHENOLATE MOFETIL
 NAFTIFINE HYDROCHLORIDE, NAFTIFINE HYDROCHLORIDE
 NALOXONE HYDROCHLORIDE, NALOXONE HYDROCHLORIDE (OTC)
 NAPROXEN, NAPROXEN
 NYSTATIN AND TRIAMCINOLONE ACETONIDE, NYSTATIN
 OLOPATADINE HYDROCHLORIDE, OLOPATADINE HYDROCHLORIDE
 ONGENTYS, OPICAPONE
 POSACONAZOLE, POSACONAZOLE
 POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
 PREDNISOLONE SODIUM PHOSPHATE, PREDNISOLONE SODIUM PHOSPHATE
 PYRIDOSTIGMINE BROMIDE, PYRIDOSTIGMINE BROMIDE
 RISPERIDONE, RISPERIDONE
 RITONAVIR, RITONAVIR
 SIROLIMUS, SIROLIMUS
 SPIRONOLACTONE, SPIRONOLACTONE
 SUCRALFATE, SUCRALFATE
 TAVABOROLE, TAVABOROLE
 TESTOSTERONE, TESTOSTERONE
 TIGECYCLINE, TIGECYCLINE
 VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE
 ZOMIG, ZOLMITRIPTAN

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ******AMNEAL PHARM**

* AMNEAL PHARMACEUTICAL
 ACEBUTOLOL HYDROCHLORIDE, ACEBUTOLOL HYDROCHLORIDE
 BETHANECHOL CHLORIDE, BETHANECHOL CHLORIDE
 DEMECLOCYCLINE HYDROCHLORIDE, DEMECLOCYCLINE HYDROCHLORIDE
 FLECAINIDE ACETATE, FLECAINIDE ACETATE
 FOLIC ACID, FOLIC ACID
 GUANFACINE HYDROCHLORIDE, GUANFACINE HYDROCHLORIDE
 HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE
 PRIMIDONE, PRIMIDONE

AMNEAL PHARMS

* AMNEAL PHARMACEUTICALS
 ACYCLOVIR, ACYCLOVIR
 ALBUTEROL SULFATE, ALBUTEROL SULFATE
 ARIPIPIRAZOLE, ARIPIPIRAZOLE
 ATOVAQUONE, ATOVAQUONE
 CALCITRIOL, CALCITRIOL
 CETIRIZINE HYDROCHLORIDE, CETIRIZINE HYDROCHLORIDE
 CHILDREN'S CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 CHILDREN'S CETIRIZINE HYDROCHLORIDE HIVES RELIEF, CETIRIZINE HYDROCHLORIDE (OTC)
 CLINDAMYCIN PALMITATE HYDROCHLORIDE, CLINDAMYCIN PALMITATE HYDROCHLORIDE
 CLOPIDOGREL BISULFATE, CLOPIDOGREL BISULFATE
 COLCHICINE, COLCHICINE
 DICLOFENAC SODIUM AND MISOPROSTOL, DICLOFENAC SODIUM
 DICLOFENAC SODIUM, DICLOFENAC SODIUM
 DICLOFENAC SODIUM, DICLOFENAC SODIUM (OTC)
 DIVALPROEX SODIUM, DIVALPROEX SODIUM
 ENTECAVIR, ENTECAVIR
 ESCITALOPRAM OXALATE, ESCITALOPRAM OXALATE
 ESTRADIOL, ESTRADIOL
 FELBAMATE, FELBAMATE
 GABAPENTIN, GABAPENTIN
 INDOMETHACIN, INDOMETHACIN
 ITRACONAZOLE, ITRACONAZOLE
 LEVETIRACETAM, LEVETIRACETAM
 LIDOCAINE, LIDOCAINE
 LORAZEPAM, LORAZEPAM
 MEMANTINE HYDROCHLORIDE AND DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
 MEROPENEM, MEROPENEM
 METAXALONE, METAXALONE
 MOMETASONE FUROATE, MOMETASONE FUROATE
 MONTELUKAST SODIUM, MONTELUKAST SODIUM
 NIACIN, NIACIN
 NITROFURANTOIN (MONOHYDRATE/MACROCRYSTALS), NITROFURANTOIN
 NITROFURANTOIN, NITROFURANTOIN
 NORETHINDRONE ACETATE, NORETHINDRONE ACETATE
 ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
 OXCARBAZEPINE, OXCARBAZEPINE
 OXYBUTYNYN CHLORIDE, OXYBUTYNYN CHLORIDE
 OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
 PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
 POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
 PROMETHAZINE HYDROCHLORIDE AND CODEINE PHOSPHATE, CODEINE PHOSPHATE
 PROMETHAZINE HYDROCHLORIDE AND DEXTROMETHORPHAN HYDROBROMIDE, DEXTROMETHORPHAN
 PROMETHAZINE HYDROCHLORIDE AND PHENYLEPHRINE HYDROCHLORIDE, PHENYLEPHRINE
 PROMETHAZINE HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE W/CODEINE PHOSPHATE, CODEINE
 QUININE SULFATE, QUININE SULFATE
 RABEPRAZOLE SODIUM, RABEPRAZOLE SODIUM
 RALOXIFENE HYDROCHLORIDE, RALOXIFENE HYDROCHLORIDE
 RISPERIDONE, RISPERIDONE
 SILDENAFIL CITRATE, SILDENAFIL CITRATE
 TELMISARTAN, TELMISARTAN
 TEMOZOLOMIDE, TEMOZOLOMIDE
 TRAMADOL HYDROCHLORIDE AND ACETAMINOPHEN, ACETAMINOPHEN
 VORICONAZOLE, VORICONAZOLE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

* AMNEAL PHARMACEUTICALS
 WARFARIN SODIUM, WARFARIN SODIUM

* AMNEAL PHARMACEUTICALS HOLDINGS GMBH
 DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE

* AMNEAL PHARMACEUTICALS OF NEW YORK LLC
 ABIRATERONE ACETATE, ABIRATERONE ACETATE
 ALOSETRON HYDROCHLORIDE, ALOSETRON HYDROCHLORIDE
 AMPHETAMINE SULFATE, AMPHETAMINE SULFATE
 ASPIRIN AND DIPYRIDAMOLE, ASPIRIN
 BENAZEPRIL HYDROCHLORIDE, BENAZEPRIL HYDROCHLORIDE
 BUDESONIDE, BUDESONIDE
 BUPRENORPHINE HYDROCHLORIDE AND NALOXONE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE
 DUTASTERIDE, DUTASTERIDE
 ERYTHROMYCIN ETHYLSUCCINATE, ERYTHROMYCIN ETHYLSUCCINATE
 GUAIFENESIN AND DEXTROMETHORPHAN HYDROBROMIDE, DEXTROMETHORPHAN HYDROBROMIDE (OTC)
 GUAIFENESIN, GUAIFENESIN (OTC)
 IBUPROFEN, IBUPROFEN (OTC)
 IRBESARTAN, IRBESARTAN
 LAMOTRIGINE, LAMOTRIGINE
 LEVONORGESTREL AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
 MECLIZINE HYDROCHLORIDE, MECLIZINE HYDROCHLORIDE
 MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
 METHYLERGONOVINE MALEATE, METHYLERGONOVINE MALEATE
 MONTELUKAST SODIUM, MONTELUKAST SODIUM
 NORETHINDRONE AND ETHINYL ESTRADIOL AND FERROUS FUMARATE, ETHINYL ESTRADIOL
 NORGESTIMATE AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
 OMEGA-3-ACID ETHYL ESTERS, OMEGA-3-ACID ETHYL ESTERS
 OSELTAMIVIR PHOSPHATE, OSELTAMIVIR PHOSPHATE
 OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN
 PALIPERIDONE, PALIPERIDONE
 PARICALCITOL, PARICALCITOL
 PRASUGREL, PRASUGREL HYDROCHLORIDE
 RIVASTIGMINE, RIVASTIGMINE
 SILDENAFIL CITRATE, SILDENAFIL CITRATE
 SPIRONOLACTONE, SPIRONOLACTONE
 SUCRALFATE, SUCRALFATE
 TOBRAMYCIN, TOBRAMYCIN
 TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE
 VALSARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 VALSARTAN, VALSARTAN
 VIGABATRIN, VIGABATRIN

AMNEAL PHARMS CO

* AMNEAL PHARMACEUTICALS CO GMBH
 ALBUTEROL SULFATE, ALBUTEROL SULFATE
 ARGATROBAN, ARGATROBAN
 BUMETANIDE, BUMETANIDE
 CHLORPROMAZINE HYDROCHLORIDE, CHLORPROMAZINE HYDROCHLORIDE
 CHLORTHALIDONE, CHLORTHALIDONE
 CLOBAZAM, CLOBAZAM
 DESIPRAMINE HYDROCHLORIDE, DESIPRAMINE HYDROCHLORIDE
 DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE
 DOXEPIN HYDROCHLORIDE, DOXEPIN HYDROCHLORIDE
 EMTRICITABINE AND TENOFOVIR DISOPROXIL FUMARATE, EMTRICITABINE
 ERYTHROMYCIN, ERYTHROMYCIN
 ETHACRYNIC ACID, ETHACRYNIC ACID
 ETODOLAC, ETODOLAC
 EZETIMIBE AND SIMVASTATIN, EZETIMIBE
 EZETIMIBE, EZETIMIBE
 FROVATRIPTAN SUCCINATE, FROVATRIPTAN SUCCINATE
 FUROSEMIDE, FUROSEMIDE
 HYDROXYCHLOROQUINE SULFATE, HYDROXYCHLOROQUINE SULFATE
 LURASIDONE HYDROCHLORIDE, LURASIDONE HYDROCHLORIDE
 NADOLOL, NADOLOL
 NAPROXEN SODIUM AND DIPHENHYDRAMINE HYDROCHLORIDE, DIPHENHYDRAMINE HYDROCHLORIDE
 OXAPROZIN, OXAPROZIN

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

* AMNEAL PHARMACEUTICALS CO GMBH
 PARICALCITOL, PARICALCITOL
 PHYTONADIONE, PHYTONADIONE
 PREGABALIN, PREGABALIN
 SEVELAMER CARBONATE, SEVELAMER CARBONATE
 SILODOSIN, SILODOSIN
 TADALAFIL, TADALAFIL
 TRANEXAMIC ACID, TRANEXAMIC ACID
 URSODIOL, URSODIOL

AMNEAL PHARMS NY

* AMNEAL PHARMACEUTICALS NY LLC
 ACETAMINOPHEN AND CODEINE PHOSPHATE, ACETAMINOPHEN
 ALPRAZOLAM, ALPRAZOLAM
 CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 CETIRIZINE HYDROCHLORIDE HIVES RELIEF, CETIRIZINE HYDROCHLORIDE (OTC)
 CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE
 EXTENDED PHENYTOIN SODIUM, PHENYTOIN SODIUM
 GABAPENTIN, GABAPENTIN
 HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
 HYDROCODONE BITARTRATE AND IBUPROFEN, HYDROCODONE BITARTRATE
 IBUPROFEN, IBUPROFEN
 IBUPROFEN, IBUPROFEN (OTC)
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 NAPROXEN SODIUM, NAPROXEN SODIUM (OTC)
 NAPROXEN, NAPROXEN
 OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN
 PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
 SULFAMETHOXAZOLE AND TRIMETHOPRIM, SULFAMETHOXAZOLE

* AMNEAL PHARMACEUTICALS OF NY LLC
 BEXAROTENE, BEXAROTENE
 ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM (OTC)
 FESOTERODINE FUMARATE, FESOTERODINE FUMARATE
 ISOTRETINOIN, ISOTRETINOIN
 OSELTAMIVIR PHOSPHATE, OSELTAMIVIR PHOSPHATE
 PITAVASTATIN CALCIUM, PITAVASTATIN CALCIUM
 PROGESTERONE, PROGESTERONE
 SILDENAFIL CITRATE, SILDENAFIL CITRATE
 TETRACYCLINE HYDROCHLORIDE, TETRACYCLINE HYDROCHLORIDE

AMNEALS PHARMS

* AMNEALS PHARMACEUTICALS LLC
 FAMOTIDINE, FAMOTIDINE

AMPHASTAR PHARM

* AMPHASTAR PHARMACEUTICAL INC
 ENOXAPARIN SODIUM (PRESERVATIVE FREE), ENOXAPARIN SODIUM

AMPHASTAR PHARMS INC

* AMPHASTAR PHARMACEUTICALS INC
 BAQSIMI, GLUCAGON
 CORTROSYN, COSYNTROPIN
 ENOXAPARIN SODIUM, ENOXAPARIN SODIUM
 GANIRELIX ACETATE, GANIRELIX ACETATE
 GLUCAGON, GLUCAGON
 ISOPROTERENOL HYDROCHLORIDE, ISOPROTERENOL HYDROCHLORIDE
 MEDROXYPROGESTERONE ACETATE, MEDROXYPROGESTERONE ACETATE
 NEOSTIGMINE METHYLSULFATE, NEOSTIGMINE METHYLSULFATE
 REXTOVY, NALOXONE HYDROCHLORIDE
 VASOPRESSIN, VASOPRESSIN

AMRING PHARMS

* AMRING PHARMACEUTICALS INC
 ARSENIC TRIOXIDE, ARSENIC TRIOXIDE
 ISOPROTERENOL HYDROCHLORIDE, ISOPROTERENOL HYDROCHLORIDE
 LAMOTRIGINE, LAMOTRIGINE
 LATANOPROST, LATANOPROST
 LYSTEDA, TRANEXAMIC ACID
 MESALAMINE, MESALAMINE
 NEOMYCIN AND POLYMYXIN B SULFATES AND HYDROCORTISONE, HYDROCORTISONE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ******* AMRING PHARMACEUTICALS INC**

NEOSTIGMINE METHYLSULFATE, NEOSTIGMINE METHYLSULFATE
 SUCCINYLCHOLINE CHLORIDE, SUCCINYLCHOLINE CHLORIDE
 TIMOLOL MALEATE, TIMOLOL MALEATE

AMTA*** AMTA LABS LTD**

DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
 MESALAMINE, MESALAMINE
 MYCOPHENOLIC SODIUM, MYCOPHENOLIC SODIUM
 PROPRANOLOL HYDROCHLORIDE, PROPRANOLOL HYDROCHLORIDE
 TOPIRAMATE, TOPIRAMATE

ANACOR PHARMS INC*** ANACOR PHARMACEUTICALS INC**

EUCRISA, CRISABOROLE

ANBEX*** ANBEX INC**

IOSAT, POTASSIUM IODIDE (OTC)

ANBISON LAB*** ANBISON LABORATORY CO LTD**

BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
 CARBAMAZEPINE, CARBAMAZEPINE
 MONTELUKAST SODIUM, MONTELUKAST SODIUM

ANDA REPOSITORY*** ANDA REPOSITORY LLC**

CALCITRIOL, CALCITRIOL
 CEFDINIR, CEFDINIR
 CEFPODOXIME PROXETIL, CEFPODOXIME PROXETIL
 CEFTRIAZONE SODIUM, CEFTRIAZONE SODIUM
 CEFUROXIME AXETIL, CEFUROXIME AXETIL
 CEPHALEXIN, CEPHALEXIN
 CYCLOBENZAPRINE HYDROCHLORIDE, CYCLOBENZAPRINE HYDROCHLORIDE
 DIATRIZOATE MEGLUMINE AND DIATRIZOATE SODIUM, DIATRIZOATE MEGLUMINE
 ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM (OTC)
 FLAC, FLUOCINOLONE ACETONIDE
 LEVETIRACETAM, LEVETIRACETAM
 LORATADINE, LORATADINE (OTC)
 METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
 OMEPRAZOLE AND SODIUM BICARBONATE, OMEPRAZOLE
 POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
 PREGABALIN, PREGABALIN
 PRIMIDONE, PRIMIDONE

ANDAS 5 HOLDING*** ANDAS 5 HOLDING LLC**

BACLOFEN, BACLOFEN
 CAPTOPRIL, CAPTOPRIL
 LABETALOL HYDROCHLORIDE, LABETALOL HYDROCHLORIDE
 RAMELTEON, RAMELTEON

ANDOR PHARMS*** ANDOR PHARMACEUTICALS LLC**

METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE

ANDRX LABS LLC*** ANDRX LABS LLC**

FORTAMET, METFORMIN HYDROCHLORIDE

ANI PHARMS*** ANI PHARMACEUTICALS INC**

ACEBUTOLOL HYDROCHLORIDE, ACEBUTOLOL HYDROCHLORIDE
 ARIMIDEX, ANASTROZOLE
 ATACAND HCT, CANDESARTAN CILEXETIL
 ATACAND, CANDESARTAN CILEXETIL
 BACLOFEN, BACLOFEN
 BENAZEPRIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, BENAZEPRIL HYDROCHLORIDE
 BENAZEPRIL HYDROCHLORIDE, BENAZEPRIL HYDROCHLORIDE
 BEXAROTENE, BEXAROTENE
 BRETHINE, TERBUTALINE SULFATE
 CASODEX, BICALUTAMIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

* ANI PHARMACEUTICALS INC
 COLESTIPOL HYDROCHLORIDE, COLESTIPOL HYDROCHLORIDE
 CORTENEMA, HYDROCORTISONE
 DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE
 ERYTHROMYCIN ETHYLSUCCINATE, ERYTHROMYCIN ETHYLSUCCINATE
 ETODOLAC, ETODOLAC
 FELBAMATE, FELBAMATE
 FENOFIBRATE (MICRONIZED), FENOFIBRATE
 FLECAINIDE ACETATE, FLECAINIDE ACETATE
 FLUCONAZOLE, FLUCONAZOLE
 GLIPIZIDE, GLIPIZIDE
 INDAPAMIDE, INDAPAMIDE
 INDERAL LA, PROPRANOLOL HYDROCHLORIDE
 INNOPRAN XL, PROPRANOLOL HYDROCHLORIDE
 KIONEX, SODIUM POLYSTYRENE SULFONATE
 LITHOBID, LITHIUM CARBONATE
 LORAZEPAM, LORAZEPAM
 LUVOX, FLUVOXAMINE MALEATE
 MEMANTINE HYDROCHLORIDE AND DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
 METHAZOLAMIDE, METHAZOLAMIDE
 METOCLOPRAMIDE HYDROCHLORIDE, METOCLOPRAMIDE HYDROCHLORIDE
 MEXILETINE HYDROCHLORIDE, MEXILETINE HYDROCHLORIDE
 MIGLUSTAT, MIGLUSTAT
 MISOPROSTOL, MISOPROSTOL
 NEBIVOLOL HYDROCHLORIDE, NEBIVOLOL HYDROCHLORIDE
 NICARDIPINE HYDROCHLORIDE, NICARDIPINE HYDROCHLORIDE
 NILUTAMIDE, NILUTAMIDE
 OXCARBAZEPINE, OXCARBAZEPINE
 OXISTAT, OXICONAZOLE NITRATE
 OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
 PENTOXIFYLLINE, PENTOXIFYLLINE
 PINDOLOL, PINDOLOL
 POLYETHYLENE GLYCOL 3350, POLYETHYLENE GLYCOL 3350 (OTC)
 POTASSIUM CITRATE, POTASSIUM CITRATE
 PROMETHAZINE HYDROCHLORIDE AND DEXTROMETHORPHAN HYDROBROMIDE, DEXTROMETHORPHAN
 PROPAFENONE HYDROCHLORIDE, PROPAFENONE HYDROCHLORIDE
 PURIFIED CORTROPHIN GEL, CORTICOTROPIN
 PYRIDOSTIGMINE BROMIDE, PYRIDOSTIGMINE BROMIDE
 REGLAN, METOCLOPRAMIDE HYDROCHLORIDE
 TRANEXAMIC ACID, TRANEXAMIC ACID
 VALPROIC ACID, VALPROIC ACID
 VANCOCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE
 VEREGEN, SINECATECHINS

ANIMA

* ANIMA PHARMACEUTICALS PVT LTD
 TRIACIN-C, CODEINE PHOSPHATE

ANNORA

* ANNORA PHARMA PRIVATE LTD
 APREMILAST, APREMILAST
 DROXIDOPA, DROXIDOPA
 LAMIVUDINE, LAMIVUDINE

ANNORA PHARMA

* ANNORA PHARMA PRIVATE LTD
 AMINOCAPROIC ACID, AMINOCAPROIC ACID
 BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
 COLCHICINE, COLCHICINE
 DEFERASIROX, DEFERASIROX
 DIATRIZOATE MEGLUMINE AND DIATRIZOATE SODIUM, DIATRIZOATE MEGLUMINE
 DICLOFENAC POTASSIUM, DICLOFENAC POTASSIUM
 DICYCLOMINE HYDROCHLORIDE, DICYCLOMINE HYDROCHLORIDE
 DIVALPROEX SODIUM, DIVALPROEX SODIUM
 ELTROMBOPAG OLAMINE, ELTROMBOPAG OLAMINE
 ENALAPRIL MALEATE, ENALAPRIL MALEATE
 EPLERENONE, EPLERENONE
 FAMOTIDINE, FAMOTIDINE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ******* ANNORA PHARMA PRIVATE LTD**

FAMOTIDINE, FAMOTIDINE (OTC)
 GABAPENTIN, GABAPENTIN
 GLYCOPYRROLATE, GLYCOPYRROLATE
 IBUPROFEN, IBUPROFEN (OTC)
 ITRACONAZOLE, ITRACONAZOLE
 IVABRADINE HYDROCHLORIDE, IVABRADINE HYDROCHLORIDE
 LURASIDONE HYDROCHLORIDE, LURASIDONE HYDROCHLORIDE
 MESALAMINE, MESALAMINE
 MEXILETINE HYDROCHLORIDE, MEXILETINE HYDROCHLORIDE
 NABUMETONE, NABUMETONE
 NIMODIPINE, NIMODIPINE
 OXCARBAZEPINE, OXCARBAZEPINE
 PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
 PIOGLITAZONE HYDROCHLORIDE, PIOGLITAZONE HYDROCHLORIDE
 POLYETHYLENE GLYCOL 3350, POLYETHYLENE GLYCOL 3350 (OTC)
 PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
 SAPROPTERIN DIHYDROCHLORIDE, SAPROPTERIN DIHYDROCHLORIDE
 SODIUM SULFATE, POTASSIUM SULFATE AND MAGNESIUM SULFATE, MAGNESIUM SULFATE
 SOLIFENACIN SUCCINATE, SOLIFENACIN SUCCINATE
 THEOPHYLLINE, THEOPHYLLINE
 VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE
 VIGABATRIN, VIGABATRIN
 ZAFIRLUKAST, ZAFIRLUKAST
 ZILEUTON, ZILEUTON

ANTARES PHARMA INC

* ANTARES PHARMA INC
 XYOSTED (AUTOINJECTOR), TESTOSTERONE ENANTHATE

ANTIBIOTICE

* ANTIBIOTICE SA
 AMPICILLIN AND SULBACTAM, AMPICILLIN SODIUM
 AMPICILLIN SODIUM, AMPICILLIN SODIUM
 NAFCILLIN SODIUM, NAFCILLIN SODIUM

AOP ORPHAN

* AOP ORPHAN PHARMACEUTICALS AG
 RAPIBLYK, LANDIOLOL HYDROCHLORIDE

APELLIS PHARMS

* APELLIS PHARMACEUTICALS INC
 EMPAVELI, PEGCETACOPLAN
 SYFOVRE, PEGCETACOPLAN

APGDI

* ASTELLAS PHARMA GLOBAL DEVELOPMENT INC
 MYRBETRIQ GRANULES, MIRABEGRON
 MYRBETRIQ, MIRABEGRON

APIIL

* ALLERGAN PHARMACEUTICALS INTERNATIONAL LTD
 ACTONEL, RISEDRONATE SODIUM
 ATELVIA, RISEDRONATE SODIUM
 LO LOESTRIN FE, ETHINYL ESTRADIOL
 TAYTULLA, ETHINYL ESTRADIOL

APNAR PHARMA LP

* APNAR PHARMA LP
 BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
 FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 MECLIZINE HYDROCHLORIDE, MECLIZINE HYDROCHLORIDE
 PRAVASTATIN SODIUM, PRAVASTATIN SODIUM
 TERAZOSIN HYDROCHLORIDE, TERAZOSIN HYDROCHLORIDE

APOTEX

* APOTEX INC
 ABIRATERONE ACETATE, ABIRATERONE ACETATE
 ACYCLOVIR, ACYCLOVIR
 ADEFOVIR DIPIVOXIL, ADEFOVIR DIPIVOXIL
 ALENDRONATE SODIUM, ALENDRONATE SODIUM
 AMBRISENTAN, AMBRISENTAN
 AMLODIPINE BESYLATE AND ATORVASTATIN CALCIUM, AMLODIPINE BESYLATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

* APOTEX INC
ARIPIPIRAZOLE, ARIPIPIRAZOLE
ATOMOXETINE HYDROCHLORIDE, ATOMOXETINE HYDROCHLORIDE
ATOVAQUONE, ATOVAQUONE
ATROPINE SULFATE, ATROPINE SULFATE
AZELASTINE HYDROCHLORIDE ALLERGY, AZELASTINE HYDROCHLORIDE (OTC)
AZELASTINE HYDROCHLORIDE AND FLUTICASONE PROPIONATE, AZELASTINE HYDROCHLORIDE
AZELASTINE HYDROCHLORIDE CHILDREN'S ALLERGY, AZELASTINE HYDROCHLORIDE (OTC)
BENZAEPRILOL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, BENAZEPRIL HYDROCHLORIDE
BENDAMUSTINE HYDROCHLORIDE, BENDAMUSTINE HYDROCHLORIDE
BEPOTASTINE BESILATE, BEPOTASTINE BESILATE
BICALUTAMIDE, BICALUTAMIDE
BIMATOPROST, BIMATOPROST
BORTEZOMIB, BORTEZOMIB
BRIMONIDINE TARTRATE AND TIMOLOL MALEATE, BRIMONIDINE TARTRATE
BRIMONIDINE TARTRATE, BRIMONIDINE TARTRATE
BROMFENAC SODIUM, BROMFENAC SODIUM
BUSULFAN, BUSULFAN
BUTORPHANOL TARTRATE, BUTORPHANOL TARTRATE
CARBIDOPA AND LEVODOPA, CARBIDOPA
CEFUROXIME AXETIL, CEFUROXIME AXETIL
CELECOXIB, CELECOXIB
CIMETIDINE, CIMETIDINE (OTC)
CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE
CYCLOSPORINE, CYCLOSPORINE
DABIGATRAN ETEXILATE MESYLATE, DABIGATRAN ETEXILATE MESYLATE
DASATINIB, DASATINIB
DESMOPRESSIN ACETATE (NEEDS NO REFRIGERATION), DESMOPRESSIN ACETATE
DEXAMETHASONE, DEXAMETHASONE
DICLOFENAC SODIUM, DICLOFENAC SODIUM
DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
DIVALPROEX SODIUM, DIVALPROEX SODIUM
DOXAZOSIN MESYLATE, DOXAZOSIN MESYLATE
DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
DOXYCYCLINE, DOXYCYCLINE
ELIGLUSTAT TARTRATE, ELIGLUSTAT TARTRATE
EPINASTINE HYDROCHLORIDE, EPINASTINE HYDROCHLORIDE
ERLOTINIB HYDROCHLORIDE, ERLOTINIB HYDROCHLORIDE
ETODOLAC, ETODOLAC
FAMCICLOVIR, FAMCICLOVIR
FAMOTIDINE, FAMOTIDINE
FENOFIBRATE (MICRONIZED), FENOFIBRATE
FINGOLIMOD HYDROCHLORIDE, FINGOLIMOD HYDROCHLORIDE
FLUPHENAZINE HYDROCHLORIDE, FLUPHENAZINE HYDROCHLORIDE
FLUTICASONE PROPIONATE, FLUTICASONE PROPIONATE (OTC)
FLUVOXAMINE MALEATE, FLUVOXAMINE MALEATE
GABAPENTIN, GABAPENTIN
GEFITINIB, GEFITINIB
GEMFIBROZIL, GEMFIBROZIL
GLIPIZIDE, GLIPIZIDE
GLYCOPYRROLATE, GLYCOPYRROLATE
GUANFACINE HYDROCHLORIDE, GUANFACINE HYDROCHLORIDE
IBANDRONATE SODIUM, IBANDRONATE SODIUM
ICOSAPENT ETHYL, ICOSAPENT ETHYL
IMATINIB MESYLATE, IMATINIB MESYLATE
IVRA, MELPHALAN HYDROCHLORIDE
LACOSAMIDE, LACOSAMIDE
LAMIVUDINE, LAMIVUDINE
LENALIDOMIDE, LENALIDOMIDE
LEVETIRACETAM, LEVETIRACETAM
LEVOCETIRIZINE DIHYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE (OTC)
LISDEXAMFETAMINE DIMESYLATE, LISDEXAMFETAMINE DIMESYLATE
LOSARTAN POTASSIUM, LOSARTAN POTASSIUM
MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

* APOTEX INC
MIDODRINE HYDROCHLORIDE, MIDODRINE HYDROCHLORIDE
MIRABEGRON, MIRABEGRON
MODAFINIL, MODAFINIL
MOMETASONE FUROATE, MOMETASONE FUROATE
MOXIFLOXACIN HYDROCHLORIDE, MOXIFLOXACIN HYDROCHLORIDE
OLOPATADINE HYDROCHLORIDE, OLOPATADINE HYDROCHLORIDE (OTC)
OMEGA-3-ACID ETHYL ESTERS, OMEGA-3-ACID ETHYL ESTERS
OMEPRAZOLE, OMEPRAZOLE
OMEPRAZOLE, OMEPRAZOLE (OTC)
ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
OXCARBAZEPINE, OXCARBAZEPINE
PAROXETINE HYDROCHLORIDE, PAROXETINE HYDROCHLORIDE
PAXIL CR, PAROXETINE HYDROCHLORIDE
PAXIL, PAROXETINE HYDROCHLORIDE
PAZOPANIB HYDROCHLORIDE, PAZOPANIB HYDROCHLORIDE
PEMETREXED DISODIUM, PEMETREXED DISODIUM
PENICILLAMINE, PENICILLAMINE
PENTOXIFYLLINE, PENTOXIFYLLINE
PIRFENIDONE, PIRFENIDONE
POMALIDOMIDE, POMALIDOMIDE
POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
PRAVASTATIN SODIUM, PRAVASTATIN SODIUM
PREGABALIN, PREGABALIN
QUINAPRIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
REGADENOSON, REGADENOSON
RISEDRONATE SODIUM, RISEDRONATE SODIUM
SELEGILINE HYDROCHLORIDE, SELEGILINE HYDROCHLORIDE
SIROLIMUS, SIROLIMUS
SOTALOL HYDROCHLORIDE, SOTALOL HYDROCHLORIDE
TAMOXIFEN CITRATE, TAMOXIFEN CITRATE
TASIMELTEON, TASIMELTEON
TERIFLUNOMIDE, TERIFLUNOMIDE
TERIPARATIDE, TERIPARATIDE
TETRABENAZINE, TETRABENAZINE
TICLOPIDINE HYDROCHLORIDE, TICLOPIDINE HYDROCHLORIDE
TIGECYCLINE, TIGECYCLINE
TIMOLOL MALEATE, TIMOLOL MALEATE
TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE
TOLVAPTAN, TOLVAPTAN
TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE
TRANEXAMIC ACID, TRANEXAMIC ACID
TRAVOPROST, TRAVOPROST
TRAZODONE HYDROCHLORIDE, TRAZODONE HYDROCHLORIDE
TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE (OTC)
TROSPIUM CHLORIDE, TROSPIUM CHLORIDE
VALACYCLOVIR HYDROCHLORIDE, VALACYCLOVIR HYDROCHLORIDE
VARENICLINE TARTRATE, VARENICLINE TARTRATE
VILAZODONE HYDROCHLORIDE, VILAZODONE HYDROCHLORIDE
ZINC SULFATE, ZINC SULFATE
ZIPRASIDONE HYDROCHLORIDE, ZIPRASIDONE HYDROCHLORIDE
ZOLEDRONIC ACID, ZOLEDRONIC ACID
ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE

APOTEX CORP

* APOTEX CORP
CUPRIC SULFATE, CUPRIC SULFATE

APOTEX INC

* APOTEX INC
ALFUZOSIN HYDROCHLORIDE, ALFUZOSIN HYDROCHLORIDE
ALPRAZOLAM, ALPRAZOLAM
ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM
AZELASTINE HYDROCHLORIDE, AZELASTINE HYDROCHLORIDE
CALCITONIN-SALMON, CALCITONIN SALMON
CARBAMAZEPINE, CARBAMAZEPINE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

* APOTEX INC
 CARBIDOPA AND LEVODOPA, CARBIDOPA
 CEFPROZIL, CEFPROZIL
 CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 CLOPIDOGREL BISULFATE, CLOPIDOGREL BISULFATE
 GRANISETRON HYDROCHLORIDE, GRANISETRON HYDROCHLORIDE
 IBANDRONATE SODIUM, IBANDRONATE SODIUM
 IMIQUIMOD, IMIQUIMOD
 IPRATROPIUM BROMIDE, IPRATROPIUM BROMIDE
 KETOTIFEN FUMARATE, KETOTIFEN FUMARATE (OTC)
 LEVETIRACETAM, LEVETIRACETAM
 LEVOFLOXACIN, LEVOFLOXACIN
 MOEXIPRIL HYDROCHLORIDE, MOEXIPRIL HYDROCHLORIDE
 MYCOPHENOLIC SODIUM, MYCOPHENOLIC SODIUM
 OFLOXACIN, OFLOXACIN
 OLANZAPINE, OLANZAPINE
 OLOPATADINE HYDROCHLORIDE, OLOPATADINE HYDROCHLORIDE
 RISPERIDONE, RISPERIDONE
 TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE
 TRAZODONE HYDROCHLORIDE, TRAZODONE HYDROCHLORIDE
 TRIAMTERENE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE

* APOTEX INC ETOBICOKE SITE
 ACYCLOVIR, ACYCLOVIR
 BALSALAZIDE DISODIUM, BALSALAZIDE DISODIUM
 BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
 CARBAMAZEPINE, CARBAMAZEPINE
 CILOSTAZOL, CILOSTAZOL
 DESMOPRESSIN ACETATE, DESMOPRESSIN ACETATE
 ETODOLAC, ETODOLAC
 FOSINOPRIL SODIUM, FOSINOPRIL SODIUM
 LEFLUNOMIDE, LEFLUNOMIDE
 LORATADINE, LORATADINE (OTC)
 MIRTAZAPINE, MIRTAZAPINE
 SELEGILINE HYDROCHLORIDE, SELEGILINE HYDROCHLORIDE
 ZONISAMIDE, ZONISAMIDE

* APOTEX INC RICHMOND HILL
 AZELASTINE HYDROCHLORIDE, AZELASTINE HYDROCHLORIDE
 BUDESONIDE, BUDESONIDE (OTC)
 FLUTICASONE PROPIONATE, FLUTICASONE PROPIONATE
 KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE

APOTHECON

* APOTHECON INC DIV BRISTOL MYERS SQUIBB
 KENALOG-10, TRIAMCINOLONE ACETONIDE
 KENALOG-40, TRIAMCINOLONE ACETONIDE
 KENALOG-80, TRIAMCINOLONE ACETONIDE

APOZEAL PHARMS

* APOZEAL PHARMACEUTICALS INC
 CETIRIZINE HYDROCHLORIDE, CETIRIZINE HYDROCHLORIDE
 CHILDREN'S CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 CHILDREN'S CETIRIZINE HYDROCHLORIDE HIVES RELIEF, CETIRIZINE HYDROCHLORIDE (OTC)
 HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE
 LACTULOSE, LACTULOSE

APP PHARMS

* APP PHARMACEUTICALS LLC
 DIPHENHYDRAMINE HYDROCHLORIDE, DIPHENHYDRAMINE HYDROCHLORIDE

APPCO

* APPCO PHARMA LLC
 ACETAZOLAMIDE, ACETAZOLAMIDE
 AMINOCAPROIC ACID, AMINOCAPROIC ACID
 BUMETANIDE, BUMETANIDE
 CHLORPROMAZINE HYDROCHLORIDE, CHLORPROMAZINE HYDROCHLORIDE
 CHLORTHALIDONE, CHLORTHALIDONE
 CLINDAMYCIN PALMITATE HYDROCHLORIDE, CLINDAMYCIN PALMITATE HYDROCHLORIDE
 CLOMIPHENE CITRATE, CLOMIPHENE CITRATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

* APPCO PHARMA LLC
 CYPROHEPTADINE HYDROCHLORIDE, CYPROHEPTADINE HYDROCHLORIDE
 DOXEPIN HYDROCHLORIDE, DOXEPIN HYDROCHLORIDE
 GLYCOPYRROLATE, GLYCOPYRROLATE
 HYDROXYCHLOROQUINE SULFATE, HYDROXYCHLOROQUINE SULFATE
 LABETALOL HYDROCHLORIDE, LABETALOL HYDROCHLORIDE
 METRONIDAZOLE, METRONIDAZOLE
 MODAFINIL, MODAFINIL
 PERPHENAZINE, PERPHENAZINE
 PRAZOSIN HYDROCHLORIDE, PRAZOSIN HYDROCHLORIDE
 SILDENAFIL CITRATE, SILDENAFIL CITRATE
 VALGANCICLOVIR HYDROCHLORIDE, VALGANCICLOVIR HYDROCHLORIDE
 VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE

APRECIA PHARMS

* APRECIA PHARMACEUTICALS LLC
 SPRITAM, LEVETIRACETAM

AQUESTIVE

* AQUESTIVE THERAPEUTICS INC
 LIBERVANT, DIAZEPAM

ARBOR PHARMS LLC

* ARBOR PHARMACEUTICALS LLC
 SKLICE, IVERMECTIN (OTC)

ARCUTIS

* ARCUTIS BIOTHERAPEUTICS INC
 ZORYVE, ROFLUMILAST

ARDELYX INC

* ARDELYX INC
 IBSRELA, TENAPANOR HYDROCHLORIDE
 XPHOZAH, TENAPANOR HYDROCHLORIDE

AREVA PHARMS

* AREVA PHARMACEUTICALS INC
 CASPOFUNGIN ACETATE, CASPOFUNGIN ACETATE
 FLUDARABINE PHOSPHATE, FLUDARABINE PHOSPHATE
 FUROSEMIDE, FUROSEMIDE
 PAMIDRONATE DISODIUM, PAMIDRONATE DISODIUM
 TERBUTALINE SULFATE, TERBUTALINE SULFATE

ARMSTRONG PHARMS

* ARMSTRONG PHARMACEUTICALS INC
 ALBUTEROL SULFATE, ALBUTEROL SULFATE
 PRIMATENE MIST, EPINEPHRINE (OTC)

ARRAY BIOPHARMA INC

* ARRAY BIOPHARMA INC
 BRAFTOVI, ENCORAFENIB
 MEKTOVI, BINIMETINIB

ARROW INTL

* ARROW INTERNATIONAL LTD
 LENALIDOMIDE, LENALIDOMIDE

ARS PHARMS OPERATION

* ARS PHARMACEUTICALS OPERATIONS INC
 NEFFY, EPINEPHRINE

ARTHUR GRP

* ARTHUR GROUP LLC
 LINCOMYCIN HYDROCHLORIDE, LINCOMYCIN HYDROCHLORIDE
 MELPHALAN HYDROCHLORIDE, MELPHALAN HYDROCHLORIDE
 SEVELAMER CARBONATE, SEVELAMER CARBONATE

ASCEND THERAPS US

* ASCEND THERAPEUTICS US LLC
 ESTROGEL, ESTRADIOL

ASCENDIS PHARMA BONE

* ASCENDIS PHARMA BONE DISEASES AS
 YORVIPATH, PALOPEGTERIPARATIDE

ASCENT PHARMS INC

* ASCENT PHARMACEUTICALS INC
 BENZONATATE, BENZONATATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

* ASCENT PHARMACEUTICALS INC
 BEXAROTENE, BEXAROTENE
 DEXMETHYLPHENIDATE HYDROCHLORIDE, DEXMETHYLPHENIDATE HYDROCHLORIDE
 DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE
 DRONABINOL, DRONABINOL
 DUTASTERIDE, DUTASTERIDE
 FAMOTIDINE, FAMOTIDINE
 FAMOTIDINE, FAMOTIDINE (OTC)
 GABAPENTIN, GABAPENTIN
 GEMFIBROZIL, GEMFIBROZIL
 HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
 HYDROMORPHONE HYDROCHLORIDE, HYDROMORPHONE HYDROCHLORIDE
 IBUPROFEN AND FAMOTIDINE, FAMOTIDINE
 IBUPROFEN, IBUPROFEN (OTC)
 ICOSAPENT ETHYL, ICOSAPENT ETHYL
 LEVOTHYROXINE SODIUM, LEVOTHYROXINE SODIUM **
 LISDEXAMFETAMINE DIMESYLATE, LISDEXAMFETAMINE DIMESYLATE
 LISINAPRIL, LISINAPRIL
 LUBIPROSTONE, LUBIPROSTONE
 MELOXICAM, MELOXICAM
 METHADONE HYDROCHLORIDE, METHADONE HYDROCHLORIDE
 METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
 OMEGA-3-ACID ETHYL ESTERS, OMEGA-3-ACID ETHYL ESTERS
 OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN
 OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
 OXYMORPHONE HYDROCHLORIDE, OXYMORPHONE HYDROCHLORIDE
 PALIPERIDONE, PALIPERIDONE
 POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
 POTASSIUM CITRATE, POTASSIUM CITRATE
 SERTRALINE HYDROCHLORIDE, SERTRALINE HYDROCHLORIDE
 TOPIRAMATE, TOPIRAMATE
 VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE

ASPEN GLOBAL INC

* ASPEN GLOBAL INC
 CYCLESSA, DESOGESTREL

ASPIRO

* ASPIRO PHARMA LTD
 ACETAMINOPHEN, ACETAMINOPHEN
 BUPIVACAINE HYDROCHLORIDE, BUPIVACAINE HYDROCHLORIDE
 CHLORPROMAZINE HYDROCHLORIDE, CHLORPROMAZINE HYDROCHLORIDE
 DAPTOMYCIN, DAPTOMYCIN
 FOSAPREPITANT DIMEGLUMINE, FOSAPREPITANT DIMEGLUMINE
 FUROSEMIDE, FUROSEMIDE
 KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
 LACOSAMIDE, LACOSAMIDE
 LIDOCAINE HYDROCHLORIDE, LIDOCAINE HYDROCHLORIDE
 OLANZAPINE, OLANZAPINE
 PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
 PHENYLEPHRINE HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE
 POSACONAZOLE, POSACONAZOLE
 PROPOFOL, PROPOFOL
 SUCCINYLCHOLINE CHLORIDE, SUCCINYLCHOLINE CHLORIDE
 SUGAMMADEX SODIUM, SUGAMMADEX SODIUM
 THIAMINE HYDROCHLORIDE, THIAMINE HYDROCHLORIDE
 VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE
 VORICONAZOLE, VORICONAZOLE

ASSERTIO

* ASSERTIO THERAPEUTICS INC
 CAMBIA, DICLOFENAC POTASSIUM
 ZIPSOR, DICLOFENAC POTASSIUM

ASTELLAS

* ASTELLAS PHARMA US INC
 AMBISOME, AMPHOTERICIN B
 ASTAGRAF XL, TACROLIMUS
 CRESEMBA, ISAVUCONAZONIUM SULFATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

* ASTELLAS PHARMA US INC
 IZERVAY, AVACINCAPTAD PEGOL SODIUM
 LEXISCAN, REGADENOSON
 MYCAMINE, MICAFUNGIN SODIUM
 PROGRAF, TACROLIMUS
 VEOZAH, FEZOLINETANT
 VESICARE LS, SOLIFENACIN SUCCINATE
 VESICARE, SOLIFENACIN SUCCINATE
 XOSPATA, GILTERITINIB FUMARATE
 XTANDI, ENZALUTAMIDE

ASTRAZENECA

* ASTRAZENECA LP
 SYMBICORT, BUDESONIDE

* ASTRAZENECA PHARMACEUTICALS LP
 AIRSUPRA, ALBUTEROL SULFATE
 BEVESPI AEROSPHERE, FORMOTEROL FUMARATE
 BRILINTA, TICAGRELOR
 DALIRESP, ROFLUMILAST
 FASLODEX, FULVESTRANT
 IRESSA, GEFITINIB
 KOSELUGO, SELUMETINIB SULFATE
 LOKELMA, SODIUM ZIRCONIUM CYCLOSILICATE
 LYNPARZA, OLAPARIB
 NEXIUM, ESOMEPRAZOLE MAGNESIUM
 PRILOSEC OTC, OMEPRAZOLE MAGNESIUM (OTC)
 PULMICORT RESPULES, BUDESONIDE
 SEROQUEL, QUETIAPINE FUMARATE
 SYMBICORT AEROSPHERE, BUDESONIDE
 TAGRISSO, OSIMERTINIB MESYLATE
 TRUQAP, CAPIVASERTIB

* ASTRAZENECA UK LTD
 CALQUENCE, ACALABRUTINIB
 CALQUENCE, ACALABRUTINIB MALEATE
 SEROQUEL XR, QUETIAPINE FUMARATE

ASTRAZENECA AB

* ASTRAZENECA AB
 BREZTRI AEROSPHERE, BUDESONIDE
 BYDUREON BCISE, EXENATIDE SYNTHETIC
 BYETTA, EXENATIDE SYNTHETIC
 FARXIGA, DAPAGLIFLOZIN
 QTERN, DAPAGLIFLOZIN
 SYMLIN, PRAMLINTIDE ACETATE
 WAINUA (AUTOINJECTOR), EPLONTERSEN SODIUM
 XIGDUO XR, DAPAGLIFLOZIN

ASTRAZENECA LP

* ASTRAZENECA LP
 NEXIUM 24HR, ESOMEPRAZOLE MAGNESIUM (OTC)

ATHEM

* ATHEM HOLDINGS LLC
 AMANTADINE HYDROCHLORIDE, AMANTADINE HYDROCHLORIDE
 FOLIC ACID, FOLIC ACID

ATHENA

* ATHENA BIOSCIENCE LLC
 NEXICLON XR, CLONIDINE

ATLAS PHARMS LLC

* ATLAS PHARMACEUTICALS LLC
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE

ATNAHS PHARMA US

* ATNAHS PHARMA US LTD
 ANAPROX DS, NAPROXEN SODIUM
 EC-NAPROSYN, NAPROXEN
 KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
 NAPROSYN, NAPROXEN

ATON

* ATON PHARMA INC

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

* ATON PHARMA INC
 LODOSYN, CARBIDOPA

AUCTA

* AUCTA PHARMACEUTICALS INC
 DEFERASIROX, DEFERASIROX
 MOTPOLY XR, LACOSAMIDE
 OXCARBAZEPINE, OXCARBAZEPINE
 VIGADRONE, VIGABATRIN

AURINIA

* AURINIA PHARMACEUTICALS INC
 LUPKYNIS, VOCLOSPORIN

AUROBINDO

* AUROBINDO PHARMA LTD
 AMOXICILLIN, AMOXICILLIN
 CEFADROXIL, CEFADROXIL/CEFADROXIL HEMIHYDRATE
 CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE
 CLARITHROMYCIN, CLARITHROMYCIN
 DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
 LISINOPRIL AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 LISINOPRIL, LISINOPRIL
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 MIRTAZAPINE, MIRTAZAPINE
 NEVIRAPINE, NEVIRAPINE
 ZIDOVUDINE, ZIDOVUDINE

AUROBINDO PHARMA

* AUROBINDO PHARMA LTD
 ACETAMINOPHEN, ACETAMINOPHEN (OTC)
 ACYCLOVIR, ACYCLOVIR
 AFIRMELLE, ETHINYL ESTRADIOL
 ALENDRONATE SODIUM, ALENDRONATE SODIUM
 ALPRAZOLAM, ALPRAZOLAM
 AMBRISENTAN, AMBRISENTAN
 AMINOCAPROIC ACID, AMINOCAPROIC ACID
 AMLODIPINE AND OLMESARTAN MEDOXOMIL, AMLODIPINE BESYLATE
 AMLODIPINE BESYLATE AND VALSARTAN, AMLODIPINE BESYLATE
 AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
 AMLODIPINE BESYLATE, VALSARTAN AND HYDROCHLOROTHIAZIDE, AMLODIPINE BESYLATE
 AMOXICILLIN AND CLAVULANATE POTASSIUM, AMOXICILLIN
 AMPICILLIN TRIHYDRATE, AMPICILLIN/AMPICILLIN TRIHYDRATE
 ARIPIPIRAZOLE, ARIPIPIRAZOLE
 ATAZANAVIR SULFATE, ATAZANAVIR SULFATE
 ATENOLOL, ATENOLOL
 ATHENTIA NEXT, LEVONORGESTREL (OTC)
 ATOMOXETINE HYDROCHLORIDE, ATOMOXETINE HYDROCHLORIDE
 AUROVELA 1.5/30, ETHINYL ESTRADIOL
 AUROVELA 1/20, ETHINYL ESTRADIOL
 AUROVELA 24 FE, ETHINYL ESTRADIOL
 AUROVELA FE 1.5/30, ETHINYL ESTRADIOL
 AUROVELA FE 1/20, ETHINYL ESTRADIOL
 AYUNA, ETHINYL ESTRADIOL
 BENAZEPRIL HYDROCHLORIDE, BENAZEPRIL HYDROCHLORIDE
 BISOPROLOL FUMARATE, BISOPROLOL FUMARATE
 CARBIDOPA, CARBIDOPA
 CARISOPRODOL, CARISOPRODOL
 CARVEDILOL, CARVEDILOL
 CEFADROXIL, CEFADROXIL/CEFADROXIL HEMIHYDRATE
 CEFDINIR, CEFDINIR
 CEFPODOXIME PROXETIL, CEFPODOXIME PROXETIL
 CEFPROZIL, CEFPROZIL
 CEFUROXIME AXETIL, CEFUROXIME AXETIL
 CELECOXIB, CELECOXIB
 CEPHALEXIN, CEPHALEXIN
 CEVIMELINE HYDROCHLORIDE, CEVIMELINE HYDROCHLORIDE
 CHILDREN'S CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 CHILDREN'S CETIRIZINE HYDROCHLORIDE HIVES RELIEF, CETIRIZINE HYDROCHLORIDE (OTC)

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ******* AUROBINDO PHARMA LTD**

CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE
 CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE
 CLINDAMYCIN HYDROCHLORIDE, CLINDAMYCIN HYDROCHLORIDE
 CLOMIPRAMINE HYDROCHLORIDE, CLOMIPRAMINE HYDROCHLORIDE
 CLORAZEPATE DIPOTASSIUM, CLORAZEPATE DIPOTASSIUM
 CLOZAPINE, CLOZAPINE
 CYCLOBENZAPRINE HYDROCHLORIDE, CYCLOBENZAPRINE HYDROCHLORIDE
 CYONANZ, ETHINYL ESTRADIOL
 DALFAMPRIDINE, DALFAMPRIDINE
 DARIFENACIN HYDROBROMIDE, DARIFENACIN HYDROBROMIDE
 DESMOPRESSIN ACETATE, DESMOPRESSIN ACETATE
 DICLOXACILLIN SODIUM, DICLOXACILLIN SODIUM
 DIMETHYL FUMARATE, DIMETHYL FUMARATE
 DUTASTERIDE AND TAMSULOSIN HYDROCHLORIDE, DUTASTERIDE
 EFAVIRENZ, EFAVIRENZ
 EFAVIRENZ, EMTRICITABINE, AND TENOFOVIR DISOPROXIL FUMARATE, EFAVIRENZ
 ELETRIPTAN HYDROBROMIDE, ELETRIPTAN HYDROBROMIDE
 EMTRICITABINE AND TENOFOVIR DISOPROXIL FUMARATE, EMTRICITABINE
 EMZAHH, NORETHINDRONE
 ENALAPRIL MALEATE, ENALAPRIL MALEATE
 ENTECAVIR, ENTECAVIR
 ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM (OTC)
 ESTRADIOL, ESTRADIOL
 ESZOPICLONE, ESZOPICLONE
 EZETIMIBE, EZETIMIBE
 FAMOTIDINE, FAMOTIDINE (OTC)
 FENOFIBRATE, FENOFIBRATE
 FESOTERODINE FUMARATE, FESOTERODINE FUMARATE
 FEXOFENADINE HYDROCHLORIDE AND PSEUDOEPHEDRINE HYDROCHLORIDE, FEXOFENADINE
 FINASTERIDE, FINASTERIDE
 FLUCONAZOLE, FLUCONAZOLE
 FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 FOSINOPRIL SODIUM AND HYDROCHLOROTHIAZIDE, FOSINOPRIL SODIUM
 GABAPENTIN, GABAPENTIN
 GALANTAMINE HYDROBROMIDE, GALANTAMINE HYDROBROMIDE
 GLIPIZIDE, GLIPIZIDE
 GLYBURIDE AND METFORMIN HYDROCHLORIDE, GLYBURIDE
 GLYCOPYRROLATE, GLYCOPYRROLATE
 GUAIFENESIN AND DEXTROMETHORPHAN HYDROBROMIDE, DEXTROMETHORPHAN HYDROBROMIDE (OTC)
 GUAIFENESIN, GUAIFENESIN (OTC)
 HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 IBANDRONATE SODIUM, IBANDRONATE SODIUM
 IBUPROFEN AND DIPHENHYDRAMINE CITRATE, DIPHENHYDRAMINE CITRATE (OTC)
 IBUPROFEN AND PSEUDOEPHEDRINE HYDROCHLORIDE, IBUPROFEN (OTC)
 IBUPROFEN, IBUPROFEN (OTC)
 INCASSIA, NORETHINDRONE
 ISOSORBIDE MONONITRATE, ISOSORBIDE MONONITRATE
 ISOTRETINOIN, ISOTRETINOIN
 KALLIGA, DESOGESTREL
 LAMIVUDINE AND ZIDOVUDINE, LAMIVUDINE
 LAMIVUDINE, LAMIVUDINE
 LAMOTRIGINE, LAMOTRIGINE
 LEFLUNOMIDE, LEFLUNOMIDE
 LEVETIRACETAM, LEVETIRACETAM
 LEVOTHYROXINE SODIUM, LEVOTHYROXINE SODIUM **
 LO SIMPESSE, ETHINYL ESTRADIOL
 LOPERAMIDE HYDROCHLORIDE, LOPERAMIDE HYDROCHLORIDE (OTC)
 LOSARTAN POTASSIUM AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 LOSARTAN POTASSIUM, LOSARTAN POTASSIUM
 MELOXICAM, MELOXICAM
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 METOPROLOL TARTRATE, METOPROLOL TARTRATE
 MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE
 MINOXIDIL (FOR MEN), MINOXIDIL (OTC)

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

* AUROBINDO PHARMA LTD
MINOXIDIL (FOR WOMEN), MINOXIDIL (OTC)
MIRTAZAPINE, MIRTAZAPINE
MOMETASONE FUROATE, MOMETASONE FUROATE
MOMETASONE FUROATE, MOMETASONE FUROATE (OTC)
MYCOPHENOLATE MOFETIL, MYCOPHENOLATE MOFETIL
NADOLOL, NADOLOL
NAPROXEN SODIUM AND PSEUDOEPHEDRINE HYDROCHLORIDE, NAPROXEN SODIUM (OTC)
NAPROXEN, NAPROXEN
NEXESTA FE, ETHINYL ESTRADIOL
NICOTINE POLACRILEX, NICOTINE POLACRILEX (OTC)
NIFEDIPINE, NIFEDIPINE
NITROFURANTOIN (MONOHYDRATE/MACROCRYSTALS), NITROFURANTOIN
NITROFURANTOIN, NITROFURANTOIN
NITROGLYCERIN, NITROGLYCERIN
NYLIA 1/35, ETHINYL ESTRADIOL
NYLIA 7/7/7, ETHINYL ESTRADIOL
OLMESARTAN MEDOXOMIL AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
OLMESARTAN MEDOXOMIL, OLMESARTAN MEDOXOMIL
OMEPRAZOLE AND SODIUM BICARBONATE, OMEPRAZOLE (OTC)
OMEPRAZOLE MAGNESIUM, OMEPRAZOLE MAGNESIUM (OTC)
OMEPRAZOLE, OMEPRAZOLE
ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
ONDANSETRON, ONDANSETRON
PAROXETINE HYDROCHLORIDE, PAROXETINE HYDROCHLORIDE
PENICILLIN V POTASSIUM, PENICILLIN V POTASSIUM
PERINDOPRIL ERBUMINE, PERINDOPRIL ERBUMINE
PHENOXYBENZAMINE HYDROCHLORIDE, PHENOXYBENZAMINE HYDROCHLORIDE
PHENYTOIN SODIUM, PHENYTOIN SODIUM
PITAVASTATIN CALCIUM, PITAVASTATIN CALCIUM
POSACONAZOLE, POSACONAZOLE
POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
PRASUGREL, PRASUGREL HYDROCHLORIDE
PRAVASTATIN SODIUM, PRAVASTATIN SODIUM
PREGABALIN, PREGABALIN
PROPAFENONE HYDROCHLORIDE, PROPAFENONE HYDROCHLORIDE
PSEUDOEPHEDRINE HYDROCHLORIDE, PSEUDOEPHEDRINE HYDROCHLORIDE (OTC)
QUETIAPINE FUMARATE, QUETIAPINE FUMARATE
QUINAPRIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
RABEPRAZOLE SODIUM, RABEPRAZOLE SODIUM
RALOXIFENE HYDROCHLORIDE, RALOXIFENE HYDROCHLORIDE
RANOLAZINE, RANOLAZINE
RIBAVIRIN, RIBAVIRIN
RISEDRONATE SODIUM, RISEDRONATE SODIUM
RIVASTIGMINE TARTRATE, RIVASTIGMINE TARTRATE
RUFINAMIDE, RUFINAMIDE
SAXAGLIPTIN, SAXAGLIPTIN HYDROCHLORIDE
SERTRALINE HYDROCHLORIDE, SERTRALINE HYDROCHLORIDE
SEVELAMER CARBONATE, SEVELAMER CARBONATE
SIMLIYA, DESOGESTREL
SIMPESS, ETHINYL ESTRADIOL
SIMVASTATIN, SIMVASTATIN
SOLIFENACIN SUCCINATE, SOLIFENACIN SUCCINATE
SPIRONOLACTONE, SPIRONOLACTONE
SULFAMETHOXAZOLE AND TRIMETHOPRIM, SULFAMETHOXAZOLE
SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
TELMISARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
TELMISARTAN, TELMISARTAN
TENOFVIR DISOPROXIL FUMARATE, TENOFVIR DISOPROXIL FUMARATE
TERBINAFFINE HYDROCHLORIDE, TERBINAFFINE HYDROCHLORIDE
TERIFLUNOMIDE, TERIFLUNOMIDE
TOPIRAMATE, TOPIRAMATE
TORSEMIDE, TORSEMIDE
TRAMADOL HYDROCHLORIDE AND ACETAMINOPHEN, ACETAMINOPHEN
TRANDOLAPRIL, TRANDOLAPRIL

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

* AUROBINDO PHARMA LTD

TRAZODONE HYDROCHLORIDE, TRAZODONE HYDROCHLORIDE
 TRI-LO-MILI, ETHINYL ESTRADIOL
 VALACYCLOVIR HYDROCHLORIDE, VALACYCLOVIR HYDROCHLORIDE
 VALGANCICLOVIR HYDROCHLORIDE, VALGANCICLOVIR HYDROCHLORIDE
 VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE
 ZAFIRLUKAST, ZAFIRLUKAST
 ZALEPLON, ZALEPLON
 ZIPRASIDONE HYDROCHLORIDE, ZIPRASIDONE HYDROCHLORIDE
 ZOLMITRIPTAN, ZOLMITRIPTAN
 ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE

AUROBINDO PHARMA LTD

* AUROBINDO PHARMA LIMITED

DIVALPROEX SODIUM, DIVALPROEX SODIUM
 FOSINOPRIL SODIUM, FOSINOPRIL SODIUM
 GALANTAMINE HYDROBROMIDE, GALANTAMINE HYDROBROMIDE
 LEVOFLOXACIN, LEVOFLOXACIN
 VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE

* AUROBINDO PHARMA LTD

ABACAVIR SULFATE AND LAMIVUDINE, ABACAVIR SULFATE
 ABACAVIR SULFATE, ABACAVIR SULFATE
 ACETAMINOPHEN AND IBUPROFEN, ACETAMINOPHEN (OTC)
 ACETAMINOPHEN, ASPIRIN AND CAFFEINE, ACETAMINOPHEN (OTC)
 ALFUZOSIN HYDROCHLORIDE, ALFUZOSIN HYDROCHLORIDE
 AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE
 AMLODIPINE BESYLATE AND BENAZEPRIL HYDROCHLORIDE, AMLODIPINE BESYLATE
 AMOXICILLIN AND CLAVULANATE POTASSIUM, AMOXICILLIN
 AMOXICILLIN, AMOXICILLIN
 ARIPIRAZOLE, ARIPIRAZOLE
 ARMODAFINIL, ARMODAFINIL
 ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM
 AZELASTINE HYDROCHLORIDE, AZELASTINE HYDROCHLORIDE
 AZITHROMYCIN, AZITHROMYCIN
 BACLOFEN, BACLOFEN
 BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
 BUSPIRONE HYDROCHLORIDE, BUSPIRONE HYDROCHLORIDE
 CARBIDOPA AND LEVODOPA, CARBIDOPA
 CEFIXIME, CEFIXIME
 CEFPODOXIME PROXETIL, CEFPODOXIME PROXETIL
 CEPHALEXIN, CEPHALEXIN
 CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 CETIRIZINE HYDROCHLORIDE HIVES RELIEF, CETIRIZINE HYDROCHLORIDE (OTC)
 CETIRIZINE HYDROCHLORIDE, CETIRIZINE HYDROCHLORIDE (OTC)
 CHILDREN'S FEXOFENADINE HYDROCHLORIDE ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
 CHLORDIAZEPOXIDE HYDROCHLORIDE AND CLIDINIUM BROMIDE, CHLORDIAZEPOXIDE HYDROCHLORIDE
 CHLORZOXAZONE, CHLORZOXAZONE
 CIMETIDINE, CIMETIDINE
 CINACALCET HYDROCHLORIDE, CINACALCET HYDROCHLORIDE
 CLINDAMYCIN PALMITATE HYDROCHLORIDE, CLINDAMYCIN PALMITATE HYDROCHLORIDE
 CLOBAZAM, CLOBAZAM
 CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
 CLOPIDOGREL BISULFATE, CLOPIDOGREL BISULFATE
 COLCHICINE, COLCHICINE
 DARUNAVIR, DARUNAVIR
 DEFERASIROX, DEFERASIROX
 DEFLAZACORT, DEFLAZACORT
 DEUTETRABENAZINE, DEUTETRABENAZINE
 DIAZEPAM, DIAZEPAM
 DICLOFENAC POTASSIUM, DICLOFENAC POTASSIUM
 DIGOXIN, DIGOXIN
 DIVALPROEX SODIUM, DIVALPROEX SODIUM
 DOCOSANOL, DOCOSANOL (OTC)
 DOFETILDE, DOFETILIDE
 DOXEPIN HYDROCHLORIDE, DOXEPIN HYDROCHLORIDE
 DROXIDOPA, DROXIDOPA

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

* AUROBINDO PHARMA LTD
 DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE
 DUTASTERIDE, DUTASTERIDE
 EFAVIRENZ, EFAVIRENZ
 EMTRICITABINE AND TENOFOVIR DISOPROXIL FUMARATE, EMTRICITABINE
 EMTRICITABINE, EMTRICITABINE
 ENTACAPONE, ENTACAPONE
 ESCITALOPRAM OXALATE, ESCITALOPRAM OXALATE
 ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM
 ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM (OTC)
 FAMCICLOVIR, FAMCICLOVIR
 FAMOTIDINE, FAMOTIDINE
 FEBUXOSTAT, FEBUXOSTAT
 FELODIPINE, FELODIPINE
 FENOFIBRATE (MICRONIZED), FENOFIBRATE
 FENOFIBRIC ACID, CHOLINE FENOFIBRATE
 FEXOFENADINE HYDROCHLORIDE ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
 FEXOFENADINE HYDROCHLORIDE, FEXOFENADINE HYDROCHLORIDE (OTC)
 FINASTERIDE, FINASTERIDE
 FINGOLIMOD HYDROCHLORIDE, FINGOLIMOD HYDROCHLORIDE
 FLECAINIDE ACETATE, FLECAINIDE ACETATE
 FLUCYTOSINE, FLUCYTOSINE
 FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 FLUPHENAZINE HYDROCHLORIDE, FLUPHENAZINE HYDROCHLORIDE
 FLUTICASONE PROPIONATE, FLUTICASONE PROPIONATE (OTC)
 GABAPENTIN, GABAPENTIN
 GEMFIBROZIL, GEMFIBROZIL
 GLIMEPIRIDE, GLIMEPIRIDE
 GLYCOPYRROLATE, GLYCOPYRROLATE
 GUAIFENESIN AND PSEUDOEPHEDRINE HYDROCHLORIDE, GUAIFENESIN (OTC)
 HALOPERIDOL, HALOPERIDOL
 HYDROCORTISONE, HYDROCORTISONE
 HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE
 IBUPROFEN AND DIPHENHYDRAMINE HYDROCHLORIDE, DIPHENHYDRAMINE HYDROCHLORIDE (OTC)
 IBUPROFEN, IBUPROFEN
 IBUPROFEN, IBUPROFEN (OTC)
 ICLEVIA, ETHINYL ESTRADIOL
 IRBESARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 IRBESARTAN, IRBESARTAN
 LACOSAMIDE, LACOSAMIDE
 LACTULOSE, LACTULOSE
 LANSOPRAZOLE, LANSOPRAZOLE
 LIDOCAINE, LIDOCAINE
 LO-ZUMANDIMINE, DROSPIRENONE
 LOPERAMIDE HYDROCHLORIDE AND SIMETHICONE, LOPERAMIDE HYDROCHLORIDE (OTC)
 LOPERAMIDE HYDROCHLORIDE, LOPERAMIDE HYDROCHLORIDE
 LORATADINE, LORATADINE (OTC)
 LURASIDONE HYDROCHLORIDE, LURASIDONE HYDROCHLORIDE
 MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
 MESALAMINE, MESALAMINE
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 METHENAMINE HIPPURATE, METHENAMINE HIPPURATE
 METHOCARBAMOL, METHOCARBAMOL
 METRONIDAZOLE, METRONIDAZOLE
 MIDODRINE HYDROCHLORIDE, MIDODRINE HYDROCHLORIDE
 MILI, ETHINYL ESTRADIOL
 MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE
 MINOXIDIL EXTRA STRENGTH (FOR MEN), MINOXIDIL (OTC)
 MODAFINIL, MODAFINIL
 MONTELUKAST SODIUM, MONTELUKAST SODIUM
 MOXIFLOXACIN HYDROCHLORIDE, MOXIFLOXACIN HYDROCHLORIDE
 MYCOPHENOLATE MOFETIL, MYCOPHENOLATE MOFETIL
 MYCOPHENOLIC SODIUM, MYCOPHENOLIC SODIUM
 NAPROXEN SODIUM, NAPROXEN SODIUM
 NAPROXEN SODIUM, NAPROXEN SODIUM (OTC)

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

- * AUROBINDO PHARMA LTD
 NEBIVOLOL HYDROCHLORIDE, NEBIVOLOL HYDROCHLORIDE
 NIACIN, NIACIN
 NICOTINE POLACRILEX, NICOTINE POLACRILEX (OTC)
 OLANZAPINE, OLANZAPINE
 OSELTAMIVIR PHOSPHATE, OSELTAMIVIR PHOSPHATE
 PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
 PARICALCITOL, PARICALCITOL
 PHENTERMINE HYDROCHLORIDE, PHENTERMINE HYDROCHLORIDE
 PILOCARPINE HYDROCHLORIDE, PILOCARPINE HYDROCHLORIDE
 PIOGLITAZONE HYDROCHLORIDE AND METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 PIOGLITAZONE HYDROCHLORIDE, PIOGLITAZONE HYDROCHLORIDE
 POLYETHYLENE GLYCOL 3350, POLYETHYLENE GLYCOL 3350 (OTC)
 PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE
 PRAZOSIN HYDROCHLORIDE, PRAZOSIN HYDROCHLORIDE
 PREDNISOLONE, PREDNISOLONE
 PREDNISON, PREDNISON
 PROPAFENONE HYDROCHLORIDE, PROPAFENONE HYDROCHLORIDE
 PYRIMETHAMINE, PYRIMETHAMINE
 QUETIAPINE FUMARATE, QUETIAPINE FUMARATE
 QUINAPRIL HYDROCHLORIDE, QUINAPRIL HYDROCHLORIDE
 RAMELTEON, RAMELTEON
 RAMIPRIL, RAMIPRIL
 REPAGLINIDE, REPAGLINIDE
 RISEDRONATE SODIUM, RISEDRONATE SODIUM
 RISPERIDONE, RISPERIDONE
 RITONAVIR, RITONAVIR
 RIZATRIPTAN BENZOATE, RIZATRIPTAN BENZOATE
 ROFLUMILAST, ROFLUMILAST
 ROSUVASTATIN CALCIUM, ROSUVASTATIN CALCIUM
 SEVELAMER HYDROCHLORIDE, SEVELAMER HYDROCHLORIDE
 SILDENAFIL CITRATE, SILDENAFIL CITRATE
 SILODOSIN, SILODOSIN
 SUMATRIPTAN AND NAPROXEN SODIUM, NAPROXEN SODIUM
 TADALAFIL, TADALAFIL
 TAMSULOSIN HYDROCHLORIDE, TAMSULOSIN HYDROCHLORIDE
 TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE
 TOPIRAMATE, TOPIRAMATE
 TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE
 TRI-MILI, ETHINYL ESTRADIOL
 URSODIOL, URSODIOL
 VALGANCICLOVIR HYDROCHLORIDE, VALGANCICLOVIR HYDROCHLORIDE
 VALSARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 VALSARTAN, VALSARTAN
 VIGABATRIN, VIGABATRIN
 VORICONAZOLE, VORICONAZOLE
 ZONISAMIDE, ZONISAMIDE
 ZUMANDIMINE, DROSPIRENONE
- * AUROBINDO PHARMA LTD INC
 ZIDOVUDINE, ZIDOVUDINE

AUROBINDO PHARMA USA

- * AUROBINDO PHARMA USA INC
 AMANTADINE HYDROCHLORIDE, AMANTADINE HYDROCHLORIDE
 BUDESONIDE, BUDESONIDE
 BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
 CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
 CLONAZEPAM, CLONAZEPAM
 DICYCLOMINE HYDROCHLORIDE, DICYCLOMINE HYDROCHLORIDE
 DOXAZOSIN MESYLATE, DOXAZOSIN MESYLATE
 ELIMITE, PERMETHRIN
 EZETIMIBE AND SIMVASTATIN, EZETIMIBE
 GLIPIZIDE, GLIPIZIDE
 GUANFACINE HYDROCHLORIDE, GUANFACINE HYDROCHLORIDE
 HYDROXYCHLOROQUINE SULFATE, HYDROXYCHLOROQUINE SULFATE
 MAXZIDE, HYDROCHLOROTHIAZIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ******* AUROBINDO PHARMA USA INC**

MAXZIDE-25, HYDROCHLOROTHIAZIDE
 MECLIZINE HYDROCHLORIDE, MECLIZINE HYDROCHLORIDE
 NITROFURANTOIN, NITROFURANTOIN, MACROCRYSTALLINE
 PAROXETINE HYDROCHLORIDE, PAROXETINE HYDROCHLORIDE
 PINDOLOL, PINDOLOL
 PRAVASTATIN SODIUM, PRAVASTATIN SODIUM
 RASAGILINE MESYLATE, RASAGILINE MESYLATE
 SOTALOL HYDROCHLORIDE, SOTALOL HYDROCHLORIDE
 TRANEXAMIC ACID, TRANEXAMIC ACID

AUROLIFE PHARMA LLC*** AUROLIFE PHARMA LLC**

ACETAMINOPHEN AND CODEINE PHOSPHATE, ACETAMINOPHEN
 AMPHETAMINE SULFATE, AMPHETAMINE SULFATE
 BUTALBITAL, ACETAMINOPHEN AND CAFFEINE, ACETAMINOPHEN
 DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE
 DEXTROAMPHETAMINE SULFATE, DEXTROAMPHETAMINE SULFATE
 DICLOFENAC SODIUM, DICLOFENAC SODIUM
 DICLOFENAC SODIUM, DICLOFENAC SODIUM (OTC)
 HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
 HYDROCODONE BITARTRATE AND IBUPROFEN, HYDROCODONE BITARTRATE
 HYDROMORPHONE HYDROCHLORIDE, HYDROMORPHONE HYDROCHLORIDE
 LORAZEPAM, LORAZEPAM
 METHADONE HYDROCHLORIDE, METHADONE HYDROCHLORIDE
 METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
 OMEPRAZOLE AND SODIUM BICARBONATE, OMEPRAZOLE
 OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN
 OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
 OXYMORPHONE HYDROCHLORIDE, OXYMORPHONE HYDROCHLORIDE
 PHENTERMINE HYDROCHLORIDE, PHENTERMINE HYDROCHLORIDE

AUSTARPHARMA*** AUSTARPHARMA LLC**

FENOFIBRATE, FENOFIBRATE
 SOLIFENACIN SUCCINATE, SOLIFENACIN SUCCINATE

AVADEL CNS*** AVADEL CNS PHARMACEUTICALS LLC**

LUMRYZ, SODIUM OXYBATE

AVANIR PHARMS*** AVANIR PHARMACEUTICALS INC**

NUEDEXTA, DEXTROMETHORPHAN HYDROBROMIDE

AVANTHI INC*** AVANTHI INC**

CHLORPHENIRAMINE MALEATE, CHLORPHENIRAMINE MALEATE (OTC)
 DEXTROAMPHETAMINE SULFATE, DEXTROAMPHETAMINE SULFATE
 DIETHYLPROPION HYDROCHLORIDE, DIETHYLPROPION HYDROCHLORIDE
 HOMATROPINE METHYLBROMIDE AND HYDROCODONE BITARTRATE, HOMATROPINE METHYLBROMIDE
 INDOMETHACIN, INDOMETHACIN
 LOMAIRA, PHENTERMINE HYDROCHLORIDE
 OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
 OXYMORPHONE HYDROCHLORIDE, OXYMORPHONE HYDROCHLORIDE

AVEMA PHARMA*** AVEMA PHARMA SOLUTIONS**

IBUPROFEN, IBUPROFEN (OTC)

AVEO PHARMS*** AVEO PHARMACEUTICALS INC**

FOTIVDA, TIVOZANIB HYDROCHLORIDE

AVERITAS*** AVERITAS PHARMA INC**

QUTENZA, CAPSAICIN

AVET*** AVET PHARMACEUTICALS INC**

ACHROMYCIN V, TETRACYCLINE HYDROCHLORIDE
 DOXERCALCIFEROL, DOXERCALCIFEROL

AVET LIFESCIENCES

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

- * AVET LIFESCIENCES LTD
 - ACARBOSE, ACARBOSE
 - ACETAZOLAMIDE SODIUM, ACETAZOLAMIDE SODIUM
 - ADENOSINE, ADENOSINE
 - AMIKACIN SULFATE, AMIKACIN SULFATE
 - BICNU, CARMUSTINE
 - CIDOFOVIR, CIDOFOVIR
 - COLISTIMETHATE SODIUM, COLISTIMETHATE SODIUM
 - FUROSEMIDE, FUROSEMIDE
 - ISOPROTERENOL HYDROCHLORIDE, ISOPROTERENOL HYDROCHLORIDE
 - METOCLOPRAMIDE, METOCLOPRAMIDE HYDROCHLORIDE
 - PALONOSETRON HYDROCHLORIDE, PALONOSETRON HYDROCHLORIDE
 - PENTAMIDINE ISETHIONATE, PENTAMIDINE ISETHIONATE
 - PROCHLORPERAZINE EDISYLATE, PROCHLORPERAZINE EDISYLATE
 - PROPOFOL, PROPOFOL
 - TRANEXAMIC ACID, TRANEXAMIC ACID
 - ZOLEDRONIC ACID, ZOLEDRONIC ACID
- * AVET LIFESCIENCES PRIVATE LTD
 - FOSCARNET SODIUM, FOSCARNET SODIUM
 - IBUTILIDE FUMARATE, IBUTILIDE FUMARATE
 - MELPHALAN HYDROCHLORIDE, MELPHALAN HYDROCHLORIDE
 - NEOSTIGMINE METHYLSULFATE, NEOSTIGMINE METHYLSULFATE
 - PHENYLEPHRINE HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE

AVID RADIOPHARMS INC

- * AVID RADIOPHARMACEUTICALS INC
 - AMYVID, FLORBETAPIR F-18
 - TAUVID, FLORTAUCIPIR F-18

AVION PHARMS

- * AVION PHARMACEUTICALS LLC
 - BALCOLTRA, ETHINYL ESTRADIOL
 - DHIVY, CARBIDOPA
 - PONSTEL, MEFENAMIC ACID

AVONDALE PHARMS

- * AVONDALE PHARMACEUTICALS LLC
 - DIFLORASONE DIACETATE, DIFLORASONE DIACETATE
 - MELOXICAM, MELOXICAM
 - NIACOR, NIACIN

AVYXA HOLDINGS

- * AVYXA HOLDINGS LLC
 - AXTLE, PEMETREXED DIPOTASSIUM
 - CYCLOPHOSPHAMIDE, CYCLOPHOSPHAMIDE
 - DOCIVYX, DOCETAXEL
 - POSFREA, PALONOSETRON HYDROCHLORIDE

AXSOME

- * AXSOME THERAPEUTICS INC
 - AUVELITY, BUPROPION HYDROCHLORIDE

AXSOME MALTA

- * AXSOME MALTA LTD
 - SUNOSI, SOLRIAMFETOL HYDROCHLORIDE

AYANA PHARMA LTD

- * AYANA PHARMA LTD
 - DOXORUBICIN HYDROCHLORIDE (LIPOSOMAL), DOXORUBICIN HYDROCHLORIDE

AYTU

- * AYTU BIOSCIENCE INC
 - KARBINAL ER, CARBINOXAMINE MALEATE

AYTU BIOPHARMA

- * AYTU BIOPHARMA INC
 - METADATE CD, METHYLPHENIDATE HYDROCHLORIDE

AZURITY

- * AZURITY PHARMACEUTICALS INC
 - AZMIRO, TESTOSTERONE CYPIONATE
 - BIDIL, HYDRALAZINE HYDROCHLORIDE
 - DANZITEN, NILOTINIB TARTRATE
 - DEXTROAMPHETAMINE SULFATE, DEXTROAMPHETAMINE SULFATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

* AZURITY PHARMACEUTICALS INC
 EDARBI, AZILSARTAN KAMEDOXOMIL
 EDARBYCLOR, AZILSARTAN KAMEDOXOMIL
 EPANED, ENALAPRIL MALEATE
 EPRONTIA, TOPIRAMATE
 ERYTHROCIN STEARATE, ERYTHROMYCIN STEARATE
 EVEKEO, AMPHETAMINE SULFATE
 FIRVANQ KIT, VANCOMYCIN HYDROCHLORIDE
 FLEQSUVY, BACLOFEN
 GLIADEL, CARMUSTINE
 HORIZANT, GABAPENTIN ENACARBIL
 KATERZIA, AMLODIPINE BENZOATE
 KONVOMEF, OMEPRAZOLE
 METARAMINOL BITARTRATE, METARAMINOL BITARTRATE
 MYHIBBIN, MYCOPHENOLATE MOFETIL
 NYMALIZE, NIMODIPINE
 PREVDUO, GLYCOPYRROLATE
 QBRELIS, LISINAPRIL
 SOTYLIZE, SOTALOL HYDROCHLORIDE
 THYQUIDITY, LEVOTHYROXINE SODIUM
 TRIPTODUR KIT, TRIPTORELIN PAMOATE
 VERELAN PM, VERAPAMIL HYDROCHLORIDE
 VERELAN, VERAPAMIL HYDROCHLORIDE
 VIVIMUSTA, BENDAMUSTINE HYDROCHLORIDE
 XATMEP, METHOTREXATE SODIUM
 ZONISADE, ZONISAMIDE

**** B ******B BRAUN**

* B BRAUN MEDICAL INC
 ACETIC ACID 0.25% IN PLASTIC CONTAINER, ACETIC ACID, GLACIAL
 AMINO ACIDS, AMINO ACIDS
 BALANCED SALT, CALCIUM CHLORIDE
 CEFAZOLIN AND DEXTROSE, CEFAZOLIN SODIUM
 CEFEPIME AND DEXTROSE IN DUPLEX CONTAINER, CEFEPIME HYDROCHLORIDE
 CEFOXITIN AND DEXTROSE IN DUPLEX CONTAINER, CEFOXITIN SODIUM
 CEFTRIAZONE AND DEXTROSE IN DUPLEX CONTAINER, CEFTRIAZONE SODIUM
 DEXTROSE 10% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 10% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 10% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 10% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 10% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 10% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 2.5% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 2.5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 2.5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 2.5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 2.5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 2.5% IN HALF-STRENGTH LACTATED RINGER'S IN PLASTIC CONTAINER, CALCIUM
 DEXTROSE 3.3% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 5% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 5% IN LACTATED RINGER'S IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 70% IN PLASTIC CONTAINER, DEXTROSE
 GLYCINE 1.5% IN PLASTIC CONTAINER, GLYCINE
 HEPARIN SODIUM 1,000 UNITS IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, HEPARIN
 HEPARIN SODIUM 2,000 UNITS IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, HEPARIN
 HEPARIN SODIUM 20,000 UNITS IN DEXTROSE 5% IN PLASTIC CONTAINER, HEPARIN SODIUM
 HEPARIN SODIUM 25,000 UNITS IN DEXTROSE 5% IN PLASTIC CONTAINER, HEPARIN SODIUM
 ISOLYTE P IN DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
 ISOLYTE S IN PLASTIC CONTAINER, MAGNESIUM CHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** B **

* B BRAUN MEDICAL INC

ISOLYTE S PH 7.4 IN PLASTIC CONTAINER, MAGNESIUM CHLORIDE
LACTATED RINGER'S IN PLASTIC CONTAINER, CALCIUM CHLORIDE
LIDOCAINE HYDROCHLORIDE 0.2% AND DEXTROSE 5% IN PLASTIC CONTAINER, LIDOCAINE
LIDOCAINE HYDROCHLORIDE 0.4% AND DEXTROSE 5% IN PLASTIC CONTAINER, LIDOCAINE
LIDOCAINE HYDROCHLORIDE 0.8% AND DEXTROSE 5% IN PLASTIC CONTAINER, LIDOCAINE
MANNITOL 10% IN PLASTIC CONTAINER, MANNITOL
MANNITOL 15% IN PLASTIC CONTAINER, MANNITOL
MANNITOL 20% IN PLASTIC CONTAINER, MANNITOL
MANNITOL 5% IN PLASTIC CONTAINER, MANNITOL
METRO I.V. IN PLASTIC CONTAINER, METRONIDAZOLE
NUTRILIPID 20%, SOYBEAN OIL
PHYSIOLYTE IN PLASTIC CONTAINER, MAGNESIUM CHLORIDE
POTASSIUM CHLORIDE 0.037% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.2% IN PLASTIC
POTASSIUM CHLORIDE 0.037% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.45% IN PLASTIC
POTASSIUM CHLORIDE 0.037% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.9% IN PLASTIC
POTASSIUM CHLORIDE 0.037% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.11% IN PLASTIC
POTASSIUM CHLORIDE 0.037% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER,
POTASSIUM CHLORIDE 0.037% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.33% IN PLASTIC
POTASSIUM CHLORIDE 0.037% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC
POTASSIUM CHLORIDE 0.037% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,
POTASSIUM CHLORIDE 0.075% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.2% IN PLASTIC
POTASSIUM CHLORIDE 0.075% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.45% IN PLASTIC
POTASSIUM CHLORIDE 0.075% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.9% IN PLASTIC
POTASSIUM CHLORIDE 0.075% IN DEXTROSE 3.3% AND SODIUM CHLORIDE 0.3% IN PLASTIC
POTASSIUM CHLORIDE 0.075% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.11% IN PLASTIC
POTASSIUM CHLORIDE 0.075% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER,
POTASSIUM CHLORIDE 0.075% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.33% IN PLASTIC
POTASSIUM CHLORIDE 0.075% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC
POTASSIUM CHLORIDE 0.075% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,
POTASSIUM CHLORIDE 0.11% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER,
POTASSIUM CHLORIDE 0.11% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.45% IN PLASTIC
POTASSIUM CHLORIDE 0.11% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,
POTASSIUM CHLORIDE 0.11% IN DEXTROSE 3.3% AND SODIUM CHLORIDE 0.3% IN PLASTIC
POTASSIUM CHLORIDE 0.11% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER,
POTASSIUM CHLORIDE 0.11% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER,
POTASSIUM CHLORIDE 0.11% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER,
POTASSIUM CHLORIDE 0.11% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER,
POTASSIUM CHLORIDE 0.11% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,
POTASSIUM CHLORIDE 0.15% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER,
POTASSIUM CHLORIDE 0.15% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.45% IN PLASTIC
POTASSIUM CHLORIDE 0.15% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,
POTASSIUM CHLORIDE 0.15% IN DEXTROSE 3.3% AND SODIUM CHLORIDE 0.3% IN PLASTIC
POTASSIUM CHLORIDE 0.15% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER,
POTASSIUM CHLORIDE 0.15% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER,
POTASSIUM CHLORIDE 0.15% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER,
POTASSIUM CHLORIDE 0.15% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER,
POTASSIUM CHLORIDE 0.15% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,
POTASSIUM CHLORIDE 0.15% IN DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE 0.15% IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, POTASSIUM
POTASSIUM CHLORIDE 0.22% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER,
POTASSIUM CHLORIDE 0.22% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.45% IN PLASTIC
POTASSIUM CHLORIDE 0.22% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,
POTASSIUM CHLORIDE 0.22% IN DEXTROSE 3.3% AND SODIUM CHLORIDE 0.3% IN PLASTIC
POTASSIUM CHLORIDE 0.22% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER,
POTASSIUM CHLORIDE 0.22% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER,
POTASSIUM CHLORIDE 0.22% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER,
POTASSIUM CHLORIDE 0.22% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER,
POTASSIUM CHLORIDE 0.22% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,
POTASSIUM CHLORIDE 0.3% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER,
POTASSIUM CHLORIDE 0.3% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER,
POTASSIUM CHLORIDE 0.3% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,
POTASSIUM CHLORIDE 0.3% IN DEXTROSE 3.3% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER,
POTASSIUM CHLORIDE 0.3% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER,
POTASSIUM CHLORIDE 0.3% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER,

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** B ******* B BRAUN MEDICAL INC**

POTASSIUM CHLORIDE 0.3% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.3% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.3% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.3% IN DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
 POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
 RINGER'S IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, SODIUM CHLORIDE
 SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, SODIUM CHLORIDE
 SODIUM CHLORIDE 0.9%, SODIUM CHLORIDE
 SODIUM CHLORIDE 3% IN PLASTIC CONTAINER, SODIUM CHLORIDE
 SODIUM CHLORIDE 5% IN PLASTIC CONTAINER, SODIUM CHLORIDE
 SORBITOL 3.3% IN PLASTIC CONTAINER, SORBITOL
 STERILE WATER FOR INJECTION IN PLASTIC CONTAINER, STERILE WATER FOR INJECTION
 STERILE WATER IN PLASTIC CONTAINER, STERILE WATER FOR IRRIGATION
 TROPHAMINE 10%, AMINO ACIDS
 TROPHAMINE, AMINO ACIDS

B BRAUN MEDICAL INC*** B BRAUN MEDICAL INC**

ACETAMINOPHEN, ACETAMINOPHEN
 BUPIVACAINE HYDROCHLORIDE, BUPIVACAINE HYDROCHLORIDE
 CALCIUM GLUCONATE, CALCIUM GLUCONATE
 HEPARIN SODIUM, HEPARIN SODIUM
 LEVETIRACETAM IN SODIUM CHLORIDE, LEVETIRACETAM
 LIDOCAINE HYDROCHLORIDE AND EPINEPHRINE, EPINEPHRINE
 LIDOCAINE HYDROCHLORIDE, LIDOCAINE HYDROCHLORIDE
 MAGNESIUM SULFATE IN DEXTROSE 5% IN PLASTIC CONTAINER, MAGNESIUM SULFATE
 MAGNESIUM SULFATE, MAGNESIUM SULFATE
 MEROPENEM AND SODIUM CHLORIDE IN DUPLEX CONTAINER, MEROPENEM
 TROMETHAMINE, TROMETHAMINE

BAJAJ*** BAJAJ MEDICAL**

CHLORHEXIDINE GLUCONATE, CHLORHEXIDINE GLUCONATE
 LACTULOSE, LACTULOSE

*** BAJAJ MEDICAL LLC**

CETIRIZINE HYDROCHLORIDE, CETIRIZINE HYDROCHLORIDE
 CHILDREN'S CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 CHILDREN'S CETIRIZINE HYDROCHLORIDE HIVES RELIEF, CETIRIZINE HYDROCHLORIDE (OTC)
 CHLORHEXIDINE GLUCONATE, CHLORHEXIDINE GLUCONATE (OTC)

BAMF*** BAMF HEALTH INC**

SODIUM FLUORIDE F-18, SODIUM FLUORIDE F-18

BANNER LIFE SCIENCES*** BANNER LIFE SCIENCES LLC**

BAFIERTAM, MONOMETHYL FUMARATE

BARR*** BARR LABORATORIES INC**

AMILORIDE HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, AMILORIDE HYDROCHLORIDE
 ARANELLE, ETHINYL ESTRADIOL
 ASPIRIN AND DIPYRIDAMOLE, ASPIRIN
 BALZIVA-28, ETHINYL ESTRADIOL
 CHLORDIAZEPOXIDE HYDROCHLORIDE, CHLORDIAZEPOXIDE HYDROCHLORIDE
 CLONAZEPAM, CLONAZEPAM
 DANAZOL, DANAZOL
 DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE
 DEXTROAMPHETAMINE SULFATE, DEXTROAMPHETAMINE SULFATE
 DIPYRIDAMOLE, DIPYRIDAMOLE
 DUTASTERIDE, DUTASTERIDE
 ESTRADIOL AND NORETHINDRONE ACETATE, ESTRADIOL
 FLUDROCORTISONE ACETATE, FLUDROCORTISONE ACETATE
 GALANTAMINE HYDROBROMIDE, GALANTAMINE HYDROBROMIDE
 HYDROXYUREA, HYDROXYUREA
 HYDROXYZINE PAMOATE, HYDROXYZINE PAMOATE
 JUNEL 1.5/30, ETHINYL ESTRADIOL
 JUNEL 1/20, ETHINYL ESTRADIOL

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** B ****

- * BARR LABORATORIES INC
 - JUNEL FE 1.5/30, ETHINYL ESTRADIOL
 - JUNEL FE 1/20, ETHINYL ESTRADIOL
 - KARIVA, DESOGESTREL
 - KELNOR, ETHINYL ESTRADIOL
 - LESSINA-28, ETHINYL ESTRADIOL
 - MEDROXYPROGESTERONE ACETATE, MEDROXYPROGESTERONE ACETATE
 - MEFLOQUINE HYDROCHLORIDE, MEFLOQUINE HYDROCHLORIDE
 - MEGESTROL ACETATE, MEGESTROL ACETATE
 - METHOTREXATE SODIUM, METHOTREXATE SODIUM
 - NALTREXONE HYDROCHLORIDE, NALTREXONE HYDROCHLORIDE
 - NIACIN, NIACIN
 - NORETHINDRONE ACETATE, NORETHINDRONE ACETATE
 - NORETHINDRONE AND ETHINYL ESTRADIOL AND FERROUS FUMARATE, ETHINYL ESTRADIOL
 - NORTREL 0.5/35-28, ETHINYL ESTRADIOL
 - NORTREL 1/35-21, ETHINYL ESTRADIOL
 - NORTREL 1/35-28, ETHINYL ESTRADIOL
 - NORTREL 7/7/7, ETHINYL ESTRADIOL
 - PORTIA-28, ETHINYL ESTRADIOL
 - SPRINTEC, ETHINYL ESTRADIOL
 - TREXALL, METHOTREXATE SODIUM
 - TRI-LEGEST FE, ETHINYL ESTRADIOL
 - TRI-SPRINTEC, ETHINYL ESTRADIOL
- * BARR LABORATORIES INC A WHOLLY OWNED SUB OF TEVA PHARMACEUTICALS USA INC
 - LANTHANUM CARBONATE, LANTHANUM CARBONATE
- * BARR PHARMACEUTICALS
 - LEUCOVORIN CALCIUM, LEUCOVORIN CALCIUM

BARR LABS INC

- * BARR LABORATORIES INC
 - ACITRETIN, ACITRETIN
 - CLOZAPINE, CLOZAPINE
 - ESTRADIOL, ESTRADIOL
 - NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL AND FERROUS FUMARATE, ETHINYL ESTRADIOL
 - NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
 - OLANZAPINE, OLANZAPINE
 - OLOPATADINE HYDROCHLORIDE, OLOPATADINE HYDROCHLORIDE (OTC)
 - TRETINOIN, TRETINOIN
 - TRI LO SPRINTEC, ETHINYL ESTRADIOL

BAUSCH

- * BAUSCH HEALTH AMERICAS INC
 - ACANYA, BENZOYL PEROXIDE
 - BRYHALI, HALOBETASOL PROPIONATE
 - DUOBRII, HALOBETASOL PROPIONATE
 - EDECIN, ETHACRYNATE SODIUM
 - EDECIN, ETHACRYNIC ACID
 - JUBLIA, EFINACONAZOLE
 - LOCOID, HYDROCORTISONE BUTYRATE
 - ONEXTON, BENZOYL PEROXIDE
 - OXSORALEN-ULTRA, METHOXSALEN
 - SYPRINE, TRIENTINE HYDROCHLORIDE
- * BAUSCH HEALTH IRELAND LTD
 - TARGRETIN, BEXAROTENE
- * BAUSCH HEALTH US LLC
 - ANCOBON, FLUCYTOSINE
 - APLENZIN, BUPROPION HYDROBROMIDE
 - ARAZLO, TAZAROTENE
 - ATIVAN, LORAZEPAM
 - CABTREO, ADAPALENE
 - CARDIZEM CD, DILTIAZEM HYDROCHLORIDE
 - CARDIZEM LA, DILTIAZEM HYDROCHLORIDE
 - CARDIZEM, DILTIAZEM HYDROCHLORIDE
 - CLINDAGEL, CLINDAMYCIN PHOSPHATE
 - DEMSER, METYROSINE
 - DESMOPRESSIN ACETATE, DESMOPRESSIN ACETATE
 - DIASSTAT ACUDIAL, DIAZEPAM

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** B ******* BAUSCH HEALTH US LLC**

DIASTAT, DIAZEPAM
 ELIDEL, PIMECROLIMUS
 FLUNISOLIDE, FLUNISOLIDE
 GRIS-PEG, GRISEOFULVIN, ULTRAMICROSIZE
 IPRATROPIUM BROMIDE, IPRATROPIUM BROMIDE
 ISORDIL, ISOSORBIDE DINITRATE
 KLARON, SULFACETAMIDE SODIUM
 LIBRAX, CHLORDIAZEPOXIDE HYDROCHLORIDE
 LIBRIUM, CHLORDIAZEPOXIDE HYDROCHLORIDE
 LOCOID, HYDROCORTISONE BUTYRATE
 LOPROX, CICLOPIROX
 LUZU, LULICONAZOLE
 MESTINON, PYRIDOSTIGMINE BROMIDE
 METROGEL-VAGINAL, METRONIDAZOLE
 MIGRANAL, DIHYDROERGOTAMINE MESYLATE
 MINOCIN, MINOCYCLINE HYDROCHLORIDE
 NORITATE, METRONIDAZOLE
 RETIN-A MICRO, TRETINOIN
 RETIN-A, TRETINOIN
 RETIN-A-MICRO, TRETINOIN
 TASMAR, TOLCAPONE
 TIAZAC, DILTIAZEM HYDROCHLORIDE
 VANOS, FLUOCINONIDE
 VASERETIC, ENALAPRIL MALEATE
 VASOTEC, ENALAPRIL MALEATE
 VIRAZOLE, RIBAVIRIN
 WELLBUTRIN XL, BUPROPION HYDROCHLORIDE
 XENAZINE, TETRABENAZINE
 XERESE, ACYCLOVIR
 ZELAPAR, SELEGILINE HYDROCHLORIDE
 ZIANA, CLINDAMYCIN PHOSPHATE
 ZOVIRAX, ACYCLOVIR
 ZYCLARA, IMIQUIMOD

BAUSCH AND LOMB*** BAUSCH AND LOMB INC**

ALAWAY, KETOTIFEN FUMARATE (OTC)
 ALREX, LOTEPIREDNOL ETABONATE
 BESIVANCE, BESIFLOXACIN HYDROCHLORIDE
 BRINZOLAMIDE, BRINZOLAMIDE
 CHILDREN'S ALAWAY, KETOTIFEN FUMARATE (OTC)
 DICLOFENAC SODIUM, DICLOFENAC SODIUM
 DORZOLAMIDE HYDROCHLORIDE AND TIMOLOL MALEATE, DORZOLAMIDE HYDROCHLORIDE
 DORZOLAMIDE HYDROCHLORIDE, DORZOLAMIDE HYDROCHLORIDE
 FLURBIPROFEN SODIUM, FLURBIPROFEN SODIUM
 ISTALOL, TIMOLOL MALEATE
 LATANOPROST, LATANOPROST
 LOTEMAX, LOTEPIREDNOL ETABONATE
 MIOCHOL-E, ACETYLCHOLINE CHLORIDE
 OFLOXACIN, OFLOXACIN
 OPCON-A, NAPHAZOLINE HYDROCHLORIDE (OTC)
 PROLENSA, BROMFENAC SODIUM
 RETISERT, FLUOCINOLONE ACETONIDE
 SULFACETAMIDE SODIUM AND PREDNISOLONE SODIUM PHOSPHATE, PREDNISOLONE SODIUM PHOSPHATE
 TROPICAMIDE, TROPICAMIDE
 VYZULTA, LATANOPROSTENE BUNOD
 ZIRGAN, GANCICLOVIR
 ZYLET, LOTEPIREDNOL ETABONATE

*** BAUSCH AND LOMB PHARMACEUTICALS INC**

BACITRACIN ZINC AND POLYMYXIN B SULFATE, BACITRACIN ZINC
 BRIMONIDINE TARTRATE, BRIMONIDINE TARTRATE
 CROMOLYN SODIUM, CROMOLYN SODIUM (OTC)
 DEXAMETHASONE SODIUM PHOSPHATE, DEXAMETHASONE SODIUM PHOSPHATE
 DEXASPORIN, DEXAMETHASONE
 ERYTHROMYCIN, ERYTHROMYCIN

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** B ****

* BAUSCH AND LOMB PHARMACEUTICALS INC
 GENTAMICIN SULFATE, GENTAMICIN SULFATE
 LEVOBUNOLOL HYDROCHLORIDE, LEVOBUNOLOL HYDROCHLORIDE
 NEOMYCIN AND POLYMYXIN B SULFATES AND BACITRACIN ZINC, BACITRACIN ZINC
 NEOMYCIN AND POLYMYXIN B SULFATES AND DEXAMETHASONE, DEXAMETHASONE
 NEOMYCIN AND POLYMYXIN B SULFATES AND GRAMICIDIN, GRAMICIDIN
 NEOMYCIN AND POLYMYXIN B SULFATES AND HYDROCORTISONE, HYDROCORTISONE
 NEOMYCIN AND POLYMYXIN B SULFATES, BACITRACIN ZINC AND HYDROCORTISONE, BACITRACIN ZINC
 OFLOXACIN, OFLOXACIN
 OTICAIR, HYDROCORTISONE
 PENTOLAIR, CYCLOPENTOLATE HYDROCHLORIDE
 PREDNISOLONE SODIUM PHOSPHATE, PREDNISOLONE SODIUM PHOSPHATE
 PROPARACAINE HYDROCHLORIDE, PROPARACAINE HYDROCHLORIDE
 SULFACETAMIDE SODIUM, SULFACETAMIDE SODIUM
 TOBRAMYCIN AND DEXAMETHASONE, DEXAMETHASONE
 TOBRAMYCIN, TOBRAMYCIN
 TRIMETHOPRIM SULFATE AND POLYMYXIN B SULFATE, POLYMYXIN B SULFATE
 TROPICAMIDE, TROPICAMIDE

BAUSCH AND LOMB INC

* BAUSCH AND LOMB INC
 ATROPINE SULFATE, ATROPINE SULFATE
 BEPREVE, BEPOTASTINE BESILATE
 LOTEMAX SM, LOTEPRDNOL ETABONATE
 LOTEMAX, LOTEPRDNOL ETABONATE
 LUMIFY PRESERVATIVE FREE, BRIMONIDINE TARTRATE (OTC)
 LUMIFY, BRIMONIDINE TARTRATE (OTC)
 METHAZOLAMIDE, METHAZOLAMIDE
 MIEBO, PERFLUOROHEXYLOCTANE
 TIMOPTIC IN OCUDOSE, TIMOLOL MALEATE
 TIMOPTIC, TIMOLOL MALEATE
 TIMOPTIC-XE, TIMOLOL MALEATE
 XIIDRA, LIFITEGRAST
 XIPERE, TRIAMCINOLONE ACETONIDE

BAUSCH LOMB IRELAND

* BAUSCH AND LOMB IRELAND LTD
 FLUORESCEIN SODIUM AND BENOXINATE HYDROCHLORIDE, BENOXINATE HYDROCHLORIDE
 TETRACAINE HYDROCHLORIDE, TETRACAINE HYDROCHLORIDE
 VISUDYNE, VERTEPORFIN

BAXTER HLTHCARE

* BAXTER HEALTHCARE CORP
 ACETIC ACID 0.25% IN PLASTIC CONTAINER, ACETIC ACID, GLACIAL
 AMINOACETIC ACID 1.5% IN PLASTIC CONTAINER, GLYCINE
 BACTOCILL IN PLASTIC CONTAINER, OXACILLIN SODIUM
 BREVIBLOC DOUBLE STRENGTH IN PLASTIC CONTAINER, ESMOLOL HYDROCHLORIDE
 BREVIBLOC IN PLASTIC CONTAINER, ESMOLOL HYDROCHLORIDE
 BREVIBLOC, ESMOLOL HYDROCHLORIDE
 CARDIOPLEGIC IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 CEFEPIME IN PLASTIC CONTAINER, CEFEPIME HYDROCHLORIDE
 CEFTRIAZONE IN PLASTIC CONTAINER, CEFTRIAZONE SODIUM
 CLINIMIX 2.75/10 SULFITE FREE IN DEXTROSE 10% IN PLASTIC CONTAINER, AMINO ACIDS
 CLINIMIX 2.75/25 SULFITE FREE IN DEXTROSE 25% IN PLASTIC CONTAINER, AMINO ACIDS
 CLINIMIX 2.75/5 SULFITE FREE IN DEXTROSE 5% IN PLASTIC CONTAINER, AMINO ACIDS
 CLINIMIX 4.25/10 SULFITE FREE IN DEXTROSE 10% IN PLASTIC CONTAINER, AMINO ACIDS
 CLINIMIX 4.25/20 SULFITE FREE IN DEXTROSE 20% IN PLASTIC CONTAINER, AMINO ACIDS
 CLINIMIX 4.25/25 SULFITE FREE IN DEXTROSE 25% IN PLASTIC CONTAINER, AMINO ACIDS
 CLINIMIX 4.25/5 SULFITE FREE IN DEXTROSE 5% IN PLASTIC CONTAINER, AMINO ACIDS
 CLINIMIX 5/10 SULFITE FREE IN DEXTROSE 10% IN PLASTIC CONTAINER, AMINO ACIDS
 CLINIMIX 5/15 SULFITE FREE IN DEXTROSE 15% IN PLASTIC CONTAINER, AMINO ACIDS
 CLINIMIX 5/20 SULFITE FREE IN DEXTROSE 20% IN PLASTIC CONTAINER, AMINO ACIDS
 CLINIMIX 5/25 SULFITE FREE IN DEXTROSE 25% IN PLASTIC CONTAINER, AMINO ACIDS
 CLINIMIX 5/35 SULFITE FREE IN DEXTROSE 35% IN PLASTIC CONTAINER, AMINO ACIDS
 CLINIMIX 8/14 SULFITE FREE IN DEXTROSE 14% IN PLASTIC CONTAINER, AMINO ACIDS
 CLINIMIX E 2.75/10 SULFITE FREE W/ ELECT IN DEXTROSE 10% W/ CALCIUM IN PLASTIC
 CLINIMIX E 2.75/25 SULFITE FREE W/ ELECT IN DEXTROSE 25% W/ CALCIUM IN PLASTIC
 CLINIMIX E 2.75/5 SULFITE FREE W/ ELECT IN DEXTROSE 5% W/ CALCIUM IN PLASTIC CONTAINER,

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** B **

* BAXTER HEALTHCARE CORP

CLINIMIX E 4.25/10 SULFITE FREE W/ ELECT IN DEXTROSE 10% W/ CALCIUM IN PLASTIC
 CLINIMIX E 4.25/20 SULFITE FREE W/ ELECT IN DEXTROSE 20% W/ CALCIUM IN PLASTIC
 CLINIMIX E 4.25/25 SULFITE FREE W/ ELECT IN DEXTROSE 25% W/ CALCIUM IN PLASTIC
 CLINIMIX E 4.25/5 SULFITE FREE W/ ELECT IN DEXTROSE 5% W/ CALCIUM IN PLASTIC CONTAINER,
 CLINIMIX E 5/10 SULFITE FREE W/ ELECT IN DEXTROSE 10% W/ CALCIUM IN PLASTIC CONTAINER,
 CLINIMIX E 5/15 SULFITE FREE W/ ELECT IN DEXTROSE 15% W/ CALCIUM IN PLASTIC CONTAINER,
 CLINIMIX E 5/20 SULFITE FREE W/ ELECT IN 20% DEXTROSE W/ CALCIUM IN PLASTIC CONTAINER,
 CLINIMIX E 5/25 SULFITE FREE W/ ELECT IN DEXTROSE 25% W/ CALCIUM IN PLASTIC CONTAINER,
 CLINIMIX E 5/35 SULFITE FREE W/ ELECT IN DEXTROSE 35% W/ CALCIUM IN PLASTIC CONTAINER,
 CLINISOL 15% SULFITE FREE IN PLASTIC CONTAINER, AMINO ACIDS
 CYCLOPHOSPHAMIDE, CYCLOPHOSPHAMIDE
 CYTOXAN, CYCLOPHOSPHAMIDE
 DEXTROSE 10% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 2.5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 5% AND ELECTROLYTE NO. 48 IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 5% AND POTASSIUM CHLORIDE 0.075% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 5% AND POTASSIUM CHLORIDE 0.15% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 5% AND POTASSIUM CHLORIDE 0.224% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 5% AND POTASSIUM CHLORIDE 0.3% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 5% IN SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 5% IN SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 5% IN SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 5% IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 10MEQ, DEXTROSE
 DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 15MEQ (K), DEXTROSE
 DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 20MEQ (K), DEXTROSE
 DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 20MEQ, DEXTROSE
 DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 30MEQ, DEXTROSE
 DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 40MEQ, DEXTROSE
 DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 5MEQ (K), DEXTROSE
 DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 5MEQ, DEXTROSE
 DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 10MEQ IN PLASTIC CONTAINER,
 DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 15MEQ IN PLASTIC CONTAINER,
 DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 20MEQ IN PLASTIC CONTAINER,
 DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 30MEQ IN PLASTIC CONTAINER,
 DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 40MEQ IN PLASTIC CONTAINER,
 DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 5MEQ IN PLASTIC CONTAINER,
 DEXTROSE 5%, SODIUM CHLORIDE 0.45% AND POTASSIUM CHLORIDE 20MEQ (K) IN PLASTIC
 DEXTROSE 50% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 70% IN PLASTIC CONTAINER, DEXTROSE
 DIANEAL LOW CALCIUM W/ DEXTROSE 1.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 DIANEAL LOW CALCIUM W/ DEXTROSE 2.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 DIANEAL LOW CALCIUM W/ DEXTROSE 4.25% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 DIANEAL LOW CALCIUM W/DEXTROSE 1.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 DIANEAL LOW CALCIUM W/DEXTROSE 2.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 DIANEAL LOW CALCIUM W/DEXTROSE 3.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 DIANEAL LOW CALCIUM W/DEXTROSE 4.25% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 DIANEAL PD-2 W/ DEXTROSE 1.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 DIANEAL PD-2 W/ DEXTROSE 2.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 DIANEAL PD-2 W/ DEXTROSE 4.25% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 DOBUTAMINE HYDROCHLORIDE IN DEXTROSE 5% IN PLASTIC CONTAINER, DOBUTAMINE HYDROCHLORIDE
 DOPAMINE HYDROCHLORIDE IN DEXTROSE 5% IN PLASTIC CONTAINER, DOPAMINE HYDROCHLORIDE
 EXTRANEAL, ICODEXTRIN
 FAMOTIDINE PRESERVATIVE FREE IN PLASTIC CONTAINER, FAMOTIDINE
 FLAGYL I.V. RTU IN PLASTIC CONTAINER, METRONIDAZOLE
 FLUCONAZOLE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, FLUCONAZOLE
 FORANE, ISOFLURANE
 GENTAMICIN SULFATE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, GENTAMICIN SULFATE
 HEPARIN SODIUM 1,000 UNITS AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, HEPARIN
 HEPARIN SODIUM 2,000 UNITS AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, HEPARIN
 IFEX, IFOSFAMIDE
 LACTATED RINGER'S AND DEXTROSE 5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 LACTATED RINGER'S IN PLASTIC CONTAINER, CALCIUM CHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** B **

* BAXTER HEALTHCARE CORP

LIDOCAINE HYDROCHLORIDE 0.2% AND DEXTROSE 5% IN PLASTIC CONTAINER, LIDOCAINE
 LIDOCAINE HYDROCHLORIDE 0.4% AND DEXTROSE 5% IN PLASTIC CONTAINER, LIDOCAINE
 LIDOCAINE HYDROCHLORIDE 0.8% AND DEXTROSE 5% IN PLASTIC CONTAINER, LIDOCAINE
 MESNEX, MESNA
 MILRINONE LACTATE IN DEXTROSE 5% IN PLASTIC CONTAINER, MILRINONE LACTATE
 NALLPEN IN PLASTIC CONTAINER, NAFICILLIN SODIUM
 NEXTERONE, AMIODARONE HYDROCHLORIDE
 NITROGLYCERIN IN DEXTROSE 5%, NITROGLYCERIN
 OSMITROL 10% IN WATER IN PLASTIC CONTAINER, MANNITOL
 OSMITROL 10% IN WATER, MANNITOL
 OSMITROL 15% IN WATER IN PLASTIC CONTAINER, MANNITOL
 OSMITROL 15% IN WATER, MANNITOL
 OSMITROL 20% IN WATER IN PLASTIC CONTAINER, MANNITOL
 OSMITROL 20% IN WATER, MANNITOL
 OSMITROL 5% IN WATER IN PLASTIC CONTAINER, MANNITOL
 OSMITROL 5% IN WATER, MANNITOL
 PENICILLIN G POTASSIUM IN PLASTIC CONTAINER, PENICILLIN G POTASSIUM
 PLASMA-LYTE 148 IN WATER IN PLASTIC CONTAINER, MAGNESIUM CHLORIDE
 PLASMA-LYTE A IN PLASTIC CONTAINER, MAGNESIUM CHLORIDE
 POTASSIUM CHLORIDE 0.15% IN SODIUM CHLORIDE 0.45%, POTASSIUM CHLORIDE
 POTASSIUM CHLORIDE 0.15% IN SODIUM CHLORIDE 0.9%, POTASSIUM CHLORIDE
 POTASSIUM CHLORIDE 0.3% AND SODIUM CHLORIDE 0.9%, POTASSIUM CHLORIDE
 POTASSIUM CHLORIDE 10MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 10MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 10MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 10MEQ IN PLASTIC CONTAINER, POTASSIUM CHLORIDE
 POTASSIUM CHLORIDE 15MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 20MEQ IN PLASTIC CONTAINER, POTASSIUM CHLORIDE
 POTASSIUM CHLORIDE 30MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 30MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 30MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 30MEQ IN PLASTIC CONTAINER, POTASSIUM CHLORIDE
 POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 40MEQ IN PLASTIC CONTAINER, POTASSIUM CHLORIDE
 POTASSIUM CHLORIDE 5MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 5MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 5MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,
 PREMASOL 10% IN PLASTIC CONTAINER, AMINO ACIDS
 PREMASOL 6% IN PLASTIC CONTAINER, AMINO ACIDS
 RINGER'S IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 SEVOFLURANE, SEVOFLURANE
 SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, SODIUM CHLORIDE
 SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, SODIUM CHLORIDE
 SODIUM CHLORIDE 0.9% IN STERILE PLASTIC CONTAINER, SODIUM CHLORIDE
 SODIUM CHLORIDE 3% IN PLASTIC CONTAINER, SODIUM CHLORIDE
 SODIUM CHLORIDE 5% IN PLASTIC CONTAINER, SODIUM CHLORIDE
 SORBITOL 3% IN PLASTIC CONTAINER, SORBITOL
 STERILE WATER FOR INJECTION IN PLASTIC CONTAINER, STERILE WATER FOR INJECTION
 STERILE WATER IN PLASTIC CONTAINER, STERILE WATER FOR IRRIGATION
 STERILE WATER, STERILE WATER FOR IRRIGATION
 SUPRANE, DESFLURANE
 TIS-U-SOL IN PLASTIC CONTAINER, MAGNESIUM SULFATE
 TIS-U-SOL, MAGNESIUM SULFATE
 TRAVASOL 10% IN PLASTIC CONTAINER, AMINO ACIDS
 TRAVASOL 5.5% IN PLASTIC CONTAINER, AMINO ACIDS
 TRAVASOL 8.5% IN PLASTIC CONTAINER, AMINO ACIDS
 VANCOMYCIN HYDROCHLORIDE IN PLASTIC CONTAINER, VANCOMYCIN HYDROCHLORIDE

* BAXTER HEALTHCARE INTERNATIONAL SPECIALTY THERAPIES DIV

PROSOL 20% SULFITE FREE IN PLASTIC CONTAINER, AMINO ACIDS

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** B ******BAXTER HLTHCARE CORP**

* BAXTER HEALTHCARE CORP
 ACETAMINOPHEN, ACETAMINOPHEN
 BENDAMUSTINE HYDROCHLORIDE, BENDAMUSTINE HYDROCHLORIDE
 BORTEZOMIB, BORTEZOMIB
 CEFAZOLIN IN DEXTROSE, CEFAZOLIN SODIUM
 CIPROFLOXACIN, CIPROFLOXACIN
 CLINDAMYCIN PHOSPHATE IN 0.9% SODIUM CHLORIDE, CLINDAMYCIN PHOSPHATE
 CLINDAMYCIN PHOSPHATE IN 5% DEXTROSE IN PLASTIC CONTAINER, CLINDAMYCIN PHOSPHATE
 CLINOLIPID 20%, OLIVE OIL
 CYCLOPHOSPHAMIDE, CYCLOPHOSPHAMIDE
 DAPTOMYCIN IN 0.9% SODIUM CHLORIDE, DAPTOMYCIN
 DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE
 DIHYDROERGOTAMINE MESYLATE, DIHYDROERGOTAMINE MESYLATE
 DOXIL (LIPOSOMAL), DOXORUBICIN HYDROCHLORIDE
 DOXORUBICIN HYDROCHLORIDE (LIPOSOMAL), DOXORUBICIN HYDROCHLORIDE
 EPTIFIBATIDE, EPTIFIBATIDE
 ERIBULIN MESYLATE, ERIBULIN MESYLATE
 FLUMAZENIL, FLUMAZENIL
 FOSAPREPITANT DIMEGLUMINE, FOSAPREPITANT DIMEGLUMINE
 FUROSEMIDE, FUROSEMIDE
 KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
 LABETALOL HYDROCHLORIDE, LABETALOL HYDROCHLORIDE
 LEVETIRACETAM IN SODIUM CHLORIDE, LEVETIRACETAM
 MAGNESIUM SULFATE IN DEXTROSE 5% IN PLASTIC CONTAINER, MAGNESIUM SULFATE
 MAGNESIUM SULFATE IN PLASTIC CONTAINER, MAGNESIUM SULFATE
 METOPROLOL TARTRATE, METOPROLOL TARTRATE
 METRONIDAZOLE IN PLASTIC CONTAINER, METRONIDAZOLE
 MICA FUNGIN IN SODIUM CHLORIDE 0.9%, MICA FUNGIN SODIUM
 NALOXONE HYDROCHLORIDE, NALOXONE HYDROCHLORIDE
 NOREPINEPHRINE BITARTRATE IN 5% DEXTROSE, NOREPINEPHRINE BITARTRATE
 NOREPINEPHRINE BITARTRATE, NOREPINEPHRINE BITARTRATE
 ONDANSETRON HYDROCHLORIDE PRESERVATIVE FREE, ONDANSETRON HYDROCHLORIDE
 PALONOSETRON HYDROCHLORIDE, PALONOSETRON HYDROCHLORIDE
 PANTOPRAZOLE SODIUM IN 0.9% SODIUM CHLORIDE, PANTOPRAZOLE SODIUM
 PEMETREXED DISODIUM, PEMETREXED DISODIUM
 PHOXILLUM B22K 4/0 IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 PHOXILLUM BK 4/2.5 IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 PRISMASOL B22GK 4/0 IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 PRISMASOL BGK 0/2.5 IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 PRISMASOL BGK 2/0 IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 PRISMASOL BGK 2/3.5 IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 PRISMASOL BGK 4/0/1.2 IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 PRISMASOL BGK 4/2.5 IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 PRISMASOL BK 0/0/1.2 IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 REGADENOSON, REGADENOSON
 SUCCINYLCHOLINE CHLORIDE, SUCCINYLCHOLINE CHLORIDE
 TOBRAMYCIN SULFATE, TOBRAMYCIN SULFATE
 TRANSDERM SCOP, SCOPOLAMINE
 VASOPRESSIN IN SODIUM CHLORIDE 0.9%, VASOPRESSIN
 ZOSYN IN PLASTIC CONTAINER, PIPERACILLIN SODIUM

* BAXTER HEALTHCARE CORP ANESTHESIA AND CRITICAL CARE
 PROTOPAM CHLORIDE, PRALIDOXIME CHLORIDE

BAYER

* BAYER HEALTHCARE LLC
 ALEVE, NAPROXEN SODIUM (OTC)

BAYER HEALTHCARE

* BAYER HEALTHCARE PHARMACEUTICALS INC
 LAMPIT, NIFURTIMOX
 NUBEQA, DAROLUTAMIDE
 VITRAKVI, LAROTRECTINIB SULFATE

BAYER HEALTHCARE LLC

* BAYER HEALTHCARE LLC
 CHILDREN'S CLARITIN, LORATADINE (OTC)
 CLARITIN HIVES RELIEF REDITAB, LORATADINE (OTC)

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** B ****

* BAYER HEALTHCARE LLC
 CLARITIN HIVES RELIEF, LORATADINE (OTC)
 CLARITIN REDITABS, LORATADINE (OTC)
 CLARITIN, LORATADINE (OTC)
 CLARITIN-D 24 HOUR, LORATADINE (OTC)
 CLARITIN-D, LORATADINE (OTC)
 LOTRIMIN ULTRA, BUTENAFINE HYDROCHLORIDE (OTC)
 MIRALAX, POLYETHYLENE GLYCOL 3350 (OTC)

BAYER HLTHCARE

* BAYER HEALTHCARE CONSUMER CARE
 ALEVE PM, DIPHENHYDRAMINE HYDROCHLORIDE (OTC)
 ASTEPRO ALLERGY, AZELASTINE HYDROCHLORIDE (OTC)
 CHILDREN'S ASTEPRO ALLERGY, AZELASTINE HYDROCHLORIDE (OTC)

* BAYER HEALTHCARE PHARMACEUTICALS INC
 ADEMPAS, RIOCIQUAT
 ANGELIQ, DROSPIRENONE
 BEYAZ, DROSPIRENONE
 CIPRO, CIPROFLOXACIN
 CIPRO, CIPROFLOXACIN HYDROCHLORIDE
 CLIMARA PRO, ESTRADIOL
 CLIMARA, ESTRADIOL
 EOVIIST, GADOXETATE DISODIUM
 GADAVIST, GADOBUTROL
 KERENDIA, FINERENONE
 KYLEENA, LEVONORGESTREL
 MENOSTAR, ESTRADIOL
 MIRENA, LEVONORGESTREL
 NATAZIA, DIENOGEST
 NEXAVAR, SORAFENIB TOSYLATE
 SAFYRAL, DROSPIRENONE
 SKYLA, LEVONORGESTREL
 STIVARGA, REGORAFENIB
 ULTRAVIST (PHARMACY BULK), IOPROMIDE
 ULTRAVIST 300, IOPROMIDE
 ULTRAVIST 370, IOPROMIDE
 VITRAKVI, LAROTRECTINIB SULFATE
 XOFIGO, RADIUM RA-223 DICHLORIDE
 YASMIN, DROSPIRENONE
 YAZ, DROSPIRENONE

BDSI

* BIODELIVERY SCIENCES INTERNATIONAL INC
 BELBUCA, BUPRENORPHINE HYDROCHLORIDE
 SYMPROIC, NALDEMEDINE TOSYLATE

BE PHARMS

* BE PHARMACEUTICALS AG
 ARFORMOTEROL TARTRATE, ARFORMOTEROL TARTRATE
 DAPTOMYCIN, DAPTOMYCIN
 ENOXAPARIN SODIUM (PRESERVATIVE FREE), ENOXAPARIN SODIUM
 FOSAPREPITANT DIMEGLUMINE, FOSAPREPITANT DIMEGLUMINE
 HEPARIN SODIUM, HEPARIN SODIUM
 NEOSTIGMINE METHYLSULFATE, NEOSTIGMINE METHYLSULFATE
 PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
 PHENYLEPHRINE HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE
 SODIUM NITROPRUSSIDE, SODIUM NITROPRUSSIDE
 SUCCINYLCHOLINE CHLORIDE, SUCCINYLCHOLINE CHLORIDE

BEACH PRODS

* BEACH PRODUCTS INC
 CLOBETASOL PROPIONATE (EMOLLIENT), CLOBETASOL PROPIONATE

BECTON DICKINSON

* BECTON DICKINSON AND CO
 CHLORHEXIDINE GLUCONATE, CHLORHEXIDINE GLUCONATE (OTC)

BECTON DICKINSON CO

* BECTON DICKINSON AND CO
 CHLORAPREP ONE-STEP FREPP, CHLORHEXIDINE GLUCONATE (OTC)
 CHLORAPREP ONE-STEP SEPP, CHLORHEXIDINE GLUCONATE (OTC)

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** B ****

* BECTON DICKINSON AND CO
 CHLORAPREP ONE-STEP, CHLORHEXIDINE GLUCONATE (OTC)
 CHLORAPREP SINGLE SWABSTICK, CHLORHEXIDINE GLUCONATE (OTC)
 CHLORAPREP TRIPLE SWABSTICK, CHLORHEXIDINE GLUCONATE (OTC)
 CHLORAPREP WITH TINT, CHLORHEXIDINE GLUCONATE (OTC)

BEIGENE

* BEIGENE USA INC
 BRUKINSA, ZANUBRUTINIB

BEIJING

* BEIJING SCIECURE PHARMACEUTICAL CO LTD
 FLUVASTATIN SODIUM, FLUVASTATIN SODIUM

BEIJING TIDE PHARM

* BEIJING TIDE PHARMACEUTICAL CO LTD
 COLESEVELAM HYDROCHLORIDE, COLESEVELAM HYDROCHLORIDE

BEIJING YILING

* BEIJING YILING BIO-ENGINEERING AND TECHNOLOGY CO LTD
 ANASTROZOLE, ANASTROZOLE
 LETROZOLE, LETROZOLE

BELCHER

* BELCHER PHARMACEUTICALS LLC
 AMINOCAPROIC ACID, AMINOCAPROIC ACID
 CEFIXIME, CEFIXIME
 CHLORZOXAZONE, CHLORZOXAZONE
 GABAPENTIN, GABAPENTIN
 LEVETIRACETAM, LEVETIRACETAM
 MEFENAMIC ACID, MEFENAMIC ACID
 POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
 SODIUM POLYSTYRENE SULFONATE, SODIUM POLYSTYRENE SULFONATE
 TACROLIMUS, TACROLIMUS

BELCHER PHARMS

* BELCHER PHARMACEUTICALS LLC
 CEPHALEXIN, CEPHALEXIN
 DESLORATADINE, DESLORATADINE

BELOTECA

* BELOTECA INC
 DIAZEPAM, DIAZEPAM
 THIOTEPA, THIOTEPA

BESINS HLTHCARE

* BESINS HEALTHCARE IRELAND LTD
 ANDROGEL, TESTOSTERONE

BEXIMCO PHARMS USA

* BEXIMCO PHARMACEUTICALS USA INC
 BACLOFEN, BACLOFEN
 CARBIDOPA, CARBIDOPA
 CYPROHEPTADINE HYDROCHLORIDE, CYPROHEPTADINE HYDROCHLORIDE
 ELETRIPTAN HYDROBROMIDE, ELETRIPTAN HYDROBROMIDE
 FLECAINIDE ACETATE, FLECAINIDE ACETATE
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 METHOCARBAMOL, METHOCARBAMOL
 MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE
 NADOLOL, NADOLOL
 NEBIVOLOL HYDROCHLORIDE, NEBIVOLOL HYDROCHLORIDE
 OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE
 SOTALOL HYDROCHLORIDE, SOTALOL HYDROCHLORIDE

BEXIMCO USA

* BEXIMCO PHARMACEUTICALS USA INC
 CARVEDILOL, CARVEDILOL

BIOCODEX SA

* BIOCODEX SA
 DIACOMIT, STIRIPENTOL

BIOCON GENERICS

* BIOCON GENERICS INC
 TRIAMTERENE, TRIAMTERENE

BIOCON LTD

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** B ****

* BIOCON LTD
FINGOLIMOD HYDROCHLORIDE, FINGOLIMOD HYDROCHLORIDE

BIOCON PHARMA

* BIOCON PHARMA INC
LIOETHYRONINE SODIUM, LIOETHYRONINE SODIUM
MYCOPHENOLIC SODIUM, MYCOPHENOLIC SODIUM
NITROFURANTOIN, NITROFURANTOIN
OXCARBAZEPINE, OXCARBAZEPINE
PRAVASTATIN SODIUM, PRAVASTATIN SODIUM

* BIOCON PHARMA LTD
ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM
DAPTOMYCIN, DAPTOMYCIN
EVEROLIMUS, EVEROLIMUS
MICAfungin SODIUM, MICAfungin SODIUM
MYCOPHENOLIC SODIUM, MYCOPHENOLIC SODIUM
POSACONAZOLE, POSACONAZOLE
ROSUVASTATIN CALCIUM, ROSUVASTATIN CALCIUM
SACUBITRIL AND VALSARTAN, SACUBITRIL
SIMVASTATIN, SIMVASTATIN
TACROLIMUS, TACROLIMUS
TERIFLUNOMIDE, TERIFLUNOMIDE

BIOCRYST

* BIOCRYST PHARMACEUTICALS INC
ORLADEYO, BEROTRALSTAT HYDROCHLORIDE
RAPIVAB, PERAMIVIR

BIOFRONTERA

* BIOFRONTERA BIOSCIENCE GMBH
AMELUZ, AMINOLEVULINIC ACID HYDROCHLORIDE

BIOGEN IDEC

* BIOGEN IDEC INC
SPINRAZA, NUSINERSEN SODIUM

BIOGEN INC

* BIOGEN INC
TECFIDERA, DIMETHYL FUMARATE
VUMERITY, DIROXIMEL FUMARATE
ZURZUVAE, ZURANOLONE

BIOGEN MA

* BIOGEN MA INC
QALSODY, TOFERSEN

BIOLINERX LTD

* BIOLINERX LTD
APHEXDA, MOTIXAFORTIDE ACETATE

BIOMARIN PHARM

* BIOMARIN PHARMACEUTICAL INC
KUVAN, SAPROPTERIN DIHYDROCHLORIDE
VOXZOGO, VOSORITIDE

BIOMEDCL RES FDN

* BIOMEDICAL RESEARCH FOUNDATION NORTHWEST LOUISIANA
AMMONIA N 13, AMMONIA N-13
FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18
SODIUM FLUORIDE F-18, SODIUM FLUORIDE F-18

BIONPHARMA

* BIONPHARMA INC
ACETAMINOPHEN AND IBUPROFEN, ACETAMINOPHEN (OTC)
AMANTADINE HYDROCHLORIDE, AMANTADINE HYDROCHLORIDE
AMPHETAMINE SULFATE, AMPHETAMINE SULFATE
ATOVAQUONE, ATOVAQUONE
AZELASTINE HYDROCHLORIDE, AZELASTINE HYDROCHLORIDE
AZITHROMYCIN, AZITHROMYCIN
BENZONATATE, BENZONATATE
BEXAROTENE, BEXAROTENE
CALCITRIOL, CALCITRIOL
CARBAMAZEPINE, CARBAMAZEPINE
CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** B ****

* BIONPHARMA INC
 CETIRIZINE HYDROCHLORIDE HIVES RELIEF, CETIRIZINE HYDROCHLORIDE (OTC)
 CEVIMELINE HYDROCHLORIDE, CEVIMELINE HYDROCHLORIDE
 CIMETIDINE, CIMETIDINE
 CLOBAZAM, CLOBAZAM
 COLESEVELAM HYDROCHLORIDE, COLESEVELAM HYDROCHLORIDE
 DEFERASIROX, DEFERASIROX
 DEXAMETHASONE, DEXAMETHASONE
 DICLOFENAC POTASSIUM, DICLOFENAC POTASSIUM
 DICYCLOMINE HYDROCHLORIDE, DICYCLOMINE HYDROCHLORIDE
 DOFETILIDE, DOFETILIDE
 DOXYLAMINE SUCCINATE AND PYRIDOXINE HYDROCHLORIDE, DOXYLAMINE SUCCINATE
 DROXIDOPA, DROXIDOPA
 ENALAPRIL MALEATE, ENALAPRIL MALEATE
 ETHOSUXIMIDE, ETHOSUXIMIDE
 FINGOLIMOD HYDROCHLORIDE, FINGOLIMOD HYDROCHLORIDE
 FLUVOXAMINE MALEATE, FLUVOXAMINE MALEATE
 GRANISETRON HYDROCHLORIDE PRESERVATIVE FREE, GRANISETRON HYDROCHLORIDE
 GRANISETRON HYDROCHLORIDE, GRANISETRON HYDROCHLORIDE
 IBUPROFEN AND DIPHENHYDRAMINE HYDROCHLORIDE, DIPHENHYDRAMINE HYDROCHLORIDE (OTC)
 IBUPROFEN, IBUPROFEN (OTC)
 IVABRADINE HYDROCHLORIDE, IVABRADINE HYDROCHLORIDE
 KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
 LOPERAMIDE HYDROCHLORIDE AND SIMETHICONE, LOPERAMIDE HYDROCHLORIDE (OTC)
 LOPERAMIDE HYDROCHLORIDE, LOPERAMIDE HYDROCHLORIDE
 LOPERAMIDE HYDROCHLORIDE, LOPERAMIDE HYDROCHLORIDE (OTC)
 LORATADINE AND PSEUDOEPHEDRINE SULFATE, LORATADINE (OTC)
 LORATADINE, LORATADINE (OTC)
 METHIMAZOLE, METHIMAZOLE
 METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
 MIDOL LIQUID GELS, IBUPROFEN (OTC)
 NAPROXEN SODIUM, NAPROXEN SODIUM (OTC)
 NICARDIPINE HYDROCHLORIDE, NICARDIPINE HYDROCHLORIDE
 NIMODIPINE, NIMODIPINE
 OXCARBAZEPINE, OXCARBAZEPINE
 OXYBUTYNYNIN CHLORIDE, OXYBUTYNYNIN CHLORIDE
 PARICALCITOL, PARICALCITOL
 POTASSIUM CITRATE, POTASSIUM CITRATE
 PROCHLORPERAZINE MALEATE, PROCHLORPERAZINE MALEATE
 PROGESTERONE, PROGESTERONE
 RUFINAMIDE, RUFINAMIDE
 SEVELAMER CARBONATE, SEVELAMER CARBONATE
 SODIUM SULFATE, POTASSIUM SULFATE AND MAGNESIUM SULFATE, MAGNESIUM SULFATE
 TERAZOSIN HYDROCHLORIDE, TERAZOSIN HYDROCHLORIDE
 TETRABENAZINE, TETRABENAZINE
 THEOPHYLLINE, THEOPHYLLINE
 VALPROIC ACID, VALPROIC ACID
 VITAMIN D, ERGOCALCIFEROL
 ZONISAMIDE, ZONISAMIDE

BIOXCEL

* BIOXCEL THERAPEUTICS INC
 IGALMI, DEXMEDETOMIDINE HYDROCHLORIDE

BLUE EARTH

* BLUE EARTH DIAGNOSTICS LTD
 AXUMIN, FLUCICLOVINE F-18
 POSLUMA, FLOTUFOLASTAT F-18 GALLIUM

BLUEPRINT MEDICINES

* BLUEPRINT MEDICINES CORP
 AVAKIT, AVAPRITINIB

BOEHRINGER INGELHEIM

* BOEHRINGER INGELHEIM
 GILOTRIF, AFATINIB DIMALEATE
 GLYXAMBI, EMPAGLIFLOZIN
 MICARDIS HCT, HYDROCHLOROTHIAZIDE
 MICARDIS, TELMISARTAN

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** B ****

* BOEHRINGER INGELHEIM PHARMACEUTICALS INC
 APTIVUS, TIPRANAVIR
 ATROVENT HFA, IPRATROPIUM BROMIDE
 COMBIVENT RESPIMAT, ALBUTEROL SULFATE
 JARDIANCE, EMPAGLIFLOZIN
 JENTADUETO XR, LINAGLIPTIN
 JENTADUETO, LINAGLIPTIN
 OFEV, NINTEDANIB ESYLATE
 PRADAXA, DABIGATRAN ETEXILATE MESYLATE
 SPIRIVA RESPIMAT, TIOTROPIUM BROMIDE
 SPIRIVA, TIOTROPIUM BROMIDE
 STIOLTO RESPIMAT, OLODATEROL HYDROCHLORIDE
 STRIVERDI RESPIMAT, OLODATEROL HYDROCHLORIDE
 SYNJARDY XR, EMPAGLIFLOZIN
 SYNJARDY, EMPAGLIFLOZIN
 TRADJENTA, LINAGLIPTIN
 TRIJARDY XR, EMPAGLIFLOZIN

BOTANIX SB

* BOTANIX SB INC
 SOFDRA, SOFPIRONIUM BROMIDE

BPI LABS

* BPI LABS LLC
 ABLYSINOL, ALCOHOL
 EPINEPHRINE, EPINEPHRINE
 MELPHALAN HYDROCHLORIDE, MELPHALAN HYDROCHLORIDE
 MYCOPHENOLATE MOFETIL HYDROCHLORIDE, MYCOPHENOLATE MOFETIL HYDROCHLORIDE
 NALOXONE HYDROCHLORIDE, NALOXONE HYDROCHLORIDE
 PENTOBARBITAL SODIUM, PENTOBARBITAL SODIUM
 ZOLEDRONIC ACID, ZOLEDRONIC ACID

BRACCO

* BRACCO DIAGNOSTICS INC
 CARDIOGEN-82, RUBIDIUM CHLORIDE RB-82
 CHOLETEC, TECHNETIUM TC-99M MEBROFENIN KIT
 CYSTOGRAFIN DILUTE, DIATRIZOATE MEGLUMINE
 CYSTOGRAFIN, DIATRIZOATE MEGLUMINE
 E-Z-HD, BARIUM SULFATE
 E-Z-PAQUE, BARIUM SULFATE
 ENTERO VU 24%, BARIUM SULFATE
 GASTROGRAFIN, DIATRIZOATE MEGLUMINE
 IOMERVU, IOMEPROL
 ISOVUE-200, IOPAMIDOL
 ISOVUE-250, IOPAMIDOL
 ISOVUE-300, IOPAMIDOL
 ISOVUE-370, IOPAMIDOL
 ISOVUE-M 200, IOPAMIDOL
 ISOVUE-M 300, IOPAMIDOL
 KINEVAC, SINCALIDE
 LIQUID E-Z-PAQUE, BARIUM SULFATE
 LUMASON, SULFUR HEXAFLUORIDE LIPID-TYPE A MICROSPHERES
 MULTIHANCE MULTIPACK, GADOBENATE DIMEGLUMINE
 MULTIHANCE, GADOBENATE DIMEGLUMINE
 PROHANCE MULTIPACK, GADOTERIDOL
 PROHANCE, GADOTERIDOL
 READI-CAT 2 SMOOTHIE, BARIUM SULFATE
 READI-CAT 2, BARIUM SULFATE
 TAGITOL V, BARIUM SULFATE
 VARIBAR HONEY, BARIUM SULFATE
 VARIBAR NECTAR, BARIUM SULFATE
 VARIBAR PUDDING, BARIUM SULFATE
 VARIBAR THIN HONEY, BARIUM SULFATE
 VARIBAR THIN LIQUID, BARIUM SULFATE

BRAEBURN

* BRAEBURN INC
 BRIXADI, BUPRENORPHINE

BRAINTREE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** B ****

* BRAINTREE LABORATORIES INC
 GOLYTELY, POLYETHYLENE GLYCOL 3350
 NULYTELY, POLYETHYLENE GLYCOL 3350
 NULYTELY-FLAVORED, POLYETHYLENE GLYCOL 3350

BRAINTREE LABS

* BRAINTREE LABORATORIES INC
 SUFLAVE, MAGNESIUM SULFATE
 SUPREP BOWEL PREP KIT, MAGNESIUM SULFATE
 SUTAB, MAGNESIUM SULFATE

BRECKENRIDGE

* BRECKENRIDGE PHARMACEUTICAL INC
 AMITRIPTYLINE HYDROCHLORIDE, AMITRIPTYLINE HYDROCHLORIDE
 ASENAPINE MALEATE, ASENAPINE MALEATE
 DABIGATRAN ETEXILATE MESYLATE, DABIGATRAN ETEXILATE MESYLATE
 DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE
 ENTECAVIR, ENTECAVIR
 EPLERENONE, EPLERENONE
 EVEROLIMUS, EVEROLIMUS
 EXEMESTANE, EXEMESTANE
 FLUPHENAZINE HYDROCHLORIDE, FLUPHENAZINE HYDROCHLORIDE
 LAMIVUDINE, LAMIVUDINE
 METHADONE HYDROCHLORIDE, METHADONE HYDROCHLORIDE
 METHYLERGONOVINE MALEATE, METHYLERGONOVINE MALEATE
 NOREPINEPHRINE BITARTRATE, NOREPINEPHRINE BITARTRATE
 OMEPRAZOLE, OMEPRAZOLE
 PENICILLAMINE, PENICILLAMINE
 RIVASTIGMINE, RIVASTIGMINE
 TETRACYCLINE HYDROCHLORIDE, TETRACYCLINE HYDROCHLORIDE
 TRIAZOLAM, TRIAZOLAM
 TRIMIPRAMINE MALEATE, TRIMIPRAMINE MALEATE

BRECKENRIDGE PHARM

* BRECKENRIDGE PHARMACEUTICAL INC
 ESTRADIOL AND NORETHINDRONE ACETATE, ESTRADIOL
 METHSCOPOLAMINE BROMIDE, METHSCOPOLAMINE BROMIDE
 OXCARBAZEPINE, OXCARBAZEPINE
 TERBINAFINE HYDROCHLORIDE, TERBINAFINE HYDROCHLORIDE

BRIDGEBIO PHARMA

* BRIDGEBIO PHARMA INC
 ATTRUBY, ACORAMIDIS HYDROCHLORIDE

BRIGHAM WOMENS

* BRIGHAM AND WOMENS HOSP
 FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18

BRIGHAM WOMENS HOSP

* BRIGHAM AND WOMENS HOSP INC
 AMMONIA N 13, AMMONIA N-13

BRIGHTGENE

* BRIGHTGENE BIO-MEDICAL TECHNOLOGY CO LTD
 ENTECAVIR, ENTECAVIR
 FONDAPARINUX SODIUM, FONDAPARINUX SODIUM

BRISTOL

* BRISTOL MYERS SQUIBB CO
 AUGTYRO, REPOTRECTINIB
 CAMZYOS, MAVACAMTEN
 EVOTAZ, ATAZANAVIR SULFATE
 KRAZATI, ADAGRASIB
 ONUREG, AZACITIDINE
 POMALYST, POMALIDOMIDE
 SOTYKTU, DEUCRAVACITINIB
 ZEPOSIA, OZANIMOD HYDROCHLORIDE

BRISTOL MYERS SQUIBB

* BRISTOL MYERS SQUIBB
 AZACTAM, AZTREONAM
 BARACLUDE, ENTECAVIR
 IDHIFA, ENASIDENIB MESYLATE
 REVLIMID, LENALIDOMIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** B ****

- * BRISTOL MYERS SQUIBB CO
REYATAZ, ATAZANAVIR SULFATE
SPRYCEL, DASATINIB
- * BRISTOL MYERS SQUIBB CO PHARMACEUTICAL RESEARCH INSTITUTE
ELIQUIS, APIXABAN

BRISTOL-MYERS

- * BRISTOL-MYERS SQUIBB CO
ABRAXANE, PACLITAXEL
COBENFY, TROSPIMUM CHLORIDE
INREBIC, FEDRATINIB HYDROCHLORIDE
ISTODAX, ROMIDEPSIN
THALOMID, THALIDOMIDE
VIDAZA, AZACITIDINE

BROOKS STERISCIENCE

- * BROOKS STERISCIENCE LTD
MEROPENEM, MEROPENEM

BTG INTL

- * BTG INTERNATIONAL INC
AURLUMYN, ILOPROST
CYANOKIT, HYDROXOCOBALAMIN
THYROSAFE, POTASSIUM IODIDE (OTC)
VISTOGARD, URIDINE TRIACETATE
XURIDEN, URIDINE TRIACETATE

BWXT ITG

- * BWXT ITG CANADA INC
INDIUM IN 111 OXYQUINOLINE, INDIUM IN-111 OXYQUINOLINE

BWXT MEDCL

- * BWXT MEDICAL LTD
IOBENGUANE I-123, IOBENGUANE SULFATE I-123

**** C ******CADILA**

- * CADILA HEALTHCARE LTD
ACETAZOLAMIDE, ACETAZOLAMIDE
ACYCLOVIR, ACYCLOVIR
BISOPROLOL FUMARATE AND HYDROCHLOROTHIAZIDE, BISOPROLOL FUMARATE
DESONIDE, DESONIDE
METRONIDAZOLE, METRONIDAZOLE
MODAFINIL, MODAFINIL
RANOLAZINE, RANOLAZINE
TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE
TRIAMTERENE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
VORICONAZOLE, VORICONAZOLE

CADILA PHARMS LTD

- * CADILA PHARMACEUTICALS LTD
ACYCLOVIR, ACYCLOVIR
BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
CELECOXIB, CELECOXIB
DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
ERYTHROMYCIN ETHYLSUCCINATE, ERYTHROMYCIN ETHYLSUCCINATE
ERYTHROMYCIN, ERYTHROMYCIN
FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
FOLIC ACID, FOLIC ACID
GEMFIBROZIL, GEMFIBROZIL
GLYBURIDE, GLYBURIDE
HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
LABETALOL HYDROCHLORIDE, LABETALOL HYDROCHLORIDE
METRONIDAZOLE, METRONIDAZOLE
NATEGLINIDE, NATEGLINIDE
NEBIVOLOL HYDROCHLORIDE, NEBIVOLOL HYDROCHLORIDE
OFLOXACIN, OFLOXACIN
OLANZAPINE, OLANZAPINE
OLMESARTAN MEDOXOMIL, OLMESARTAN MEDOXOMIL
PAROXETINE HYDROCHLORIDE, PAROXETINE HYDROCHLORIDE
RALOXIFENE HYDROCHLORIDE, RALOXIFENE HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** C ******* CADILA PHARMACEUTICALS LTD**

RIVASTIGMINE TARTRATE, RIVASTIGMINE TARTRATE
 ROSUVASTATIN CALCIUM, ROSUVASTATIN CALCIUM
 SILDENAFIL CITRATE, SILDENAFIL CITRATE
 TELMISARTAN, TELMISARTAN
 TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE
 VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE
 VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE

CALL INC

* CALL INC DBA ROCHESTER PHARMACEUTICALS
 ADAPALENE, ADAPALENE

CALLIDITAS

* CALLIDITAS THERAPEUTICS AB
 TARPEYO, BUDESONIDE

CAPELLON PHARMS LLC

* CAPELLON PHARMACEUTICALS LLC
 POLMON, DEXCHLORPHENIRAMINE MALEATE

CAPLIN

* CAPLIN STERILES LTD
 ARGATROBAN, ARGATROBAN
 CARBOPROST TROMETHAMINE, CARBOPROST TROMETHAMINE
 CISATRACURIUM BESYLATE, CISATRACURIUM BESYLATE
 DIFLUPREDNATE, DIFLUPREDNATE
 EPHEDRINE SULFATE, EPHEDRINE SULFATE
 ETOMIDATE, ETOMIDATE
 GLYCOPYRROLATE, GLYCOPYRROLATE
 KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
 LABETALOL HYDROCHLORIDE, LABETALOL HYDROCHLORIDE
 MILRINONE LACTATE, MILRINONE LACTATE
 NEOSTIGMINE METHYLSULFATE, NEOSTIGMINE METHYLSULFATE
 NOREPINEPHRINE BITARTRATE, NOREPINEPHRINE BITARTRATE
 OFLOXACIN, OFLOXACIN
 PHENYLEPHRINE HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE
 PROCHLORPERAZINE EDISYLATE, PROCHLORPERAZINE EDISYLATE
 ROCURONIUM BROMIDE, ROCURONIUM BROMIDE
 ROPIVACAINE HYDROCHLORIDE, ROPIVACAINE HYDROCHLORIDE
 SODIUM NITROPRUSSIDE, SODIUM NITROPRUSSIDE
 SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
 THIAMINE HYDROCHLORIDE, THIAMINE HYDROCHLORIDE
 TIMOLOL MALEATE, TIMOLOL MALEATE
 TRANEXAMIC ACID, TRANEXAMIC ACID
 VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE

CARA THERAP

* CARA THERAPEUTICS INC
 KORSUVA, DIFELIKEFALIN ACETATE

CARDINAL HEALTH 414

* CARDINAL HEALTH 414 LLC CARDINAL HEALTH NUCLEAR PHARMACY SERVICES
 FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18
 LYMPHOSEEK KIT, TECHNETIUM TC-99M TILMANOCEPT
 SODIUM FLUORIDE F-18, SODIUM FLUORIDE F-18
 TECHNETIUM TC 99M SESTAMIBI, TECHNETIUM TC-99M SESTAMIBI KIT
 TECHNETIUM TC-99M MEDRONATE KIT, TECHNETIUM TC-99M MEDRONATE KIT

CARDINAL HEALTH 418

* CARDINAL HEALTH 418 INC
 SODIUM IODIDE I 123, SODIUM IODIDE I-123

CARDINAL HLTH 414

* CARDINAL HEALTH 414 LLC
 AMMONIA N 13, AMMONIA N-13

CARLSBAD

* CARLSBAD TECHNOLOGY INC
 ACYCLOVIR, ACYCLOVIR
 CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE
 DICLOFENAC SODIUM, DICLOFENAC SODIUM
 FAMOTIDINE, FAMOTIDINE
 GLIMEPIRIDE, GLIMEPIRIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** C ****

* CARLSBAD TECHNOLOGY INC
LOVASTATIN, LOVASTATIN

CARLSBAD TECHNOLOGY

* CARLSBAD TECHNOLOGY INC
ACYCLOVIR, ACYCLOVIR

CARNEGIE

* CARNEGIE PHARMACEUTICALS LLC
AMINOCAPROIC ACID, AMINOCAPROIC ACID
E.E.S. 400, ERYTHROMYCIN ETHYLSUCCINATE
E.E.S., ERYTHROMYCIN ETHYLSUCCINATE
ERY-TAB, ERYTHROMYCIN
ERYPED, ERYTHROMYCIN ETHYLSUCCINATE
ERYTHROMYCIN ETHYLSUCCINATE, ERYTHROMYCIN ETHYLSUCCINATE
ERYTHROMYCIN, ERYTHROMYCIN
ETRAVIRINE, ETRAVIRINE
FAMOTIDINE, FAMOTIDINE
PRIMIDONE, PRIMIDONE

CASPER PHARMA LLC

* CASPER PHARMA LLC
ANTIVERT, MECLIZINE HYDROCHLORIDE
AQUASOL A, VITAMIN A PALMITATE
CASPORYN HC, HYDROCORTISONE
EDETATE CALCIUM DISODIUM, EDETATE CALCIUM DISODIUM
FURADANTIN, NITROFURANTOIN
LUMI-SPORYN, BACITRACIN ZINC
ROBINUL FORTE, GLYCOPYRROLATE
ROBINUL, GLYCOPYRROLATE
THALITONE, CHLORTHALIDONE
ZYLOPRIM, ALLOPURINOL

CATALENT

* CATALENT PHARMA SOLUTIONS LLC
CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
NAPROXEN SODIUM, NAPROXEN SODIUM (OTC)
VALPROIC ACID, VALPROIC ACID

CATALYST PHARMS

* CATALYST PHARMACEUTICALS INC
AGAMREE, VAMOROLONE
FIRDAPSE, AMIFAMPRIDINE PHOSPHATE
FYCOMPA, PERAMPANEL

CEDIPROF INC

* CEDIPROF INC
DEXMETHYLPHENIDATE HYDROCHLORIDE, DEXMETHYLPHENIDATE HYDROCHLORIDE
LEVO-T, LEVOTHYROXINE SODIUM **
METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
OSELTAMIVIR PHOSPHATE, OSELTAMIVIR PHOSPHATE

CELATOR PHARMS

* CELATOR PHARMACEUTICALS INC
VYXEOS, CYTARABINE

CEPHALON

* CEPHALON INC
GABITRIL, TIAGABINE HYDROCHLORIDE
NUVIGIL, ARMODAFINIL
TREANDA, BENDAMUSTINE HYDROCHLORIDE
TRISENOX, ARSENIC TRIOXIDE
* CEPHALON LLC
PROVIGIL, MODAFINIL

CF PHARMTECH

* CF PHARMTECH INC
ARFORMOTEROL TARTRATE, ARFORMOTEROL TARTRATE

CHANGZHOU PHARM

* CHANGZHOU PHARMACEUTICAL FACTORY
CAPTOPRIL, CAPTOPRIL
DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
PREGABALIN, PREGABALIN

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** C ****

* CHANGZHOU PHARMACEUTICAL FACTORY
 ROSUVASTATIN CALCIUM, ROSUVASTATIN CALCIUM

CHARTWELL

* CHARTWELL LIFE MOLECULES LLC
 ALLOPURINOL, ALLOPURINOL
 AMOXICILLIN, AMOXICILLIN
 CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE
 CIPROFLOXACIN, CIPROFLOXACIN
 CLARITHROMYCIN, CLARITHROMYCIN
 COLOCORT, HYDROCORTISONE
 DUTASTERIDE, DUTASTERIDE
 FENOFIBRATE (MICRONIZED), FENOFIBRATE
 FLUCONAZOLE, FLUCONAZOLE
 NALTREXONE HYDROCHLORIDE, NALTREXONE HYDROCHLORIDE
 OMEGA-3-ACID ETHYL ESTERS, OMEGA-3-ACID ETHYL ESTERS
 OXAPROZIN, OXAPROZIN
 TEMOZOLOMIDE, TEMOZOLOMIDE
 TENOFOVIR DISOPROXIL FUMARATE, TENOFOVIR DISOPROXIL FUMARATE
 TERBINAFINE HYDROCHLORIDE, TERBINAFINE HYDROCHLORIDE

* CHARTWELL PHARMA SCIENCE LLC
 SULFASALAZINE, SULFASALAZINE

* CHARTWELL SCHEDULED LLC
 ACETAMINOPHEN AND CODEINE PHOSPHATE, ACETAMINOPHEN
 BENZPHETAMINE HYDROCHLORIDE, BENZPHETAMINE HYDROCHLORIDE
 DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
 DOXYCYCLINE, DOXYCYCLINE
 HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN
 PHENDIMETRAZINE TARTRATE, PHENDIMETRAZINE TARTRATE

CHARTWELL INJECTABLE

* CHARTWELL INJECTABLES LLC
 DIPYRIDAMOLE, DIPYRIDAMOLE
 DOXAPRAM HYDROCHLORIDE, DOXAPRAM HYDROCHLORIDE
 ENALAPRILAT, ENALAPRILAT

CHARTWELL MOLECULAR

* CHARTWELL MOLECULAR HOLDINGS LLC
 ACETAZOLAMIDE, ACETAZOLAMIDE
 ALBUTEROL SULFATE, ALBUTEROL SULFATE
 BROMPHENIRAMINE MALEATE, PSEUDOEPHEDRINE HYDROCHLORIDE AND DEXTROMETHORPHAN
 CALCIUM ACETATE, CALCIUM ACETATE
 CARVEDILOL, CARVEDILOL
 CETIRIZINE HYDROCHLORIDE, CETIRIZINE HYDROCHLORIDE
 CHILDREN'S CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 CHILDREN'S CETIRIZINE HYDROCHLORIDE HIVES RELIEF, CETIRIZINE HYDROCHLORIDE (OTC)
 CIMETIDINE HYDROCHLORIDE, CIMETIDINE HYDROCHLORIDE
 CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE
 CLINDAMYCIN HYDROCHLORIDE, CLINDAMYCIN HYDROCHLORIDE
 CLOBAZAM, CLOBAZAM
 CYPROHEPTADINE HYDROCHLORIDE, CYPROHEPTADINE HYDROCHLORIDE
 DEXAMETHASONE, DEXAMETHASONE
 DIAZEPAM, DIAZEPAM
 DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
 ESCITALOPRAM OXALATE, ESCITALOPRAM OXALATE
 FOLIC ACID, FOLIC ACID
 GLIMEPIRIDE, GLIMEPIRIDE
 GRANISETRON HYDROCHLORIDE, GRANISETRON HYDROCHLORIDE
 IRBESARTAN, IRBESARTAN
 ISOSORBIDE MONONITRATE, ISOSORBIDE MONONITRATE
 LAMIVUDINE, LAMIVUDINE
 LANSOPRAZOLE, LANSOPRAZOLE
 LEVETIRACETAM, LEVETIRACETAM
 LEVOCETIRIZINE DIHYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE
 LEVOFLOXACIN, LEVOFLOXACIN
 LIDOCAINE HYDROCHLORIDE VISCOUS, LIDOCAINE HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** C ****

* CHARTWELL MOLECULAR HOLDINGS LLC
 MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
 MONTELUKAST SODIUM, MONTELUKAST SODIUM
 OLANZAPINE, OLANZAPINE
 ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
 PREDNISOLONE, PREDNISOLONE
 PREDNISONE, PREDNISONE
 RAMIPRIL, RAMIPRIL
 RIFAMPIN, RIFAMPIN
 RISPERIDONE, RISPERIDONE
 SULFAMETHOXAZOLE AND TRIMETHOPRIM, SULFAMETHOXAZOLE
 THEOPHYLLINE, THEOPHYLLINE
 ZALEPLON, ZALEPLON
 ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE

CHARTWELL MOLECULES

* CHARTWELL MOLECULES LLC
 CIMETIDINE, CIMETIDINE
 CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
 DISULFIRAM, DISULFIRAM
 GEMFIBROZIL, GEMFIBROZIL
 INDOMETHACIN, INDOMETHACIN
 NABUMETONE, NABUMETONE
 ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
 ONDANSETRON, ONDANSETRON
 SULFAMETHOXAZOLE AND TRIMETHOPRIM, SULFAMETHOXAZOLE

CHARTWELL RX

* CHARTWELL RX SCIENCES LLC
 ABACAVIR SULFATE AND LAMIVUDINE, ABACAVIR SULFATE
 ABACAVIR SULFATE, ABACAVIR SULFATE
 ACTICLATE, DOXYCYCLINE HYCLATE
 ACYCLOVIR, ACYCLOVIR
 ALBUTEROL SULFATE, ALBUTEROL SULFATE
 ALPRAZOLAM, ALPRAZOLAM
 AMANTADINE HYDROCHLORIDE, AMANTADINE HYDROCHLORIDE
 AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE
 AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
 AMOXAPINE, AMOXAPINE
 AMOXICILLIN, AMOXICILLIN
 ARIPIPIRAZOLE, ARIPIPIRAZOLE
 ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM
 ATOVAQUONE, ATOVAQUONE
 AZITHROMYCIN, AZITHROMYCIN
 BENAZEPRIL HYDROCHLORIDE, BENAZEPRIL HYDROCHLORIDE
 BENZTROPINE MESYLATE, BENZTROPINE MESYLATE
 BETHANECHOL CHLORIDE, BETHANECHOL CHLORIDE
 BICALUTAMIDE, BICALUTAMIDE
 CALCIPOTRIENE, CALCIPOTRIENE
 CALCIUM ACETATE, CALCIUM ACETATE
 CARBAMAZEPINE, CARBAMAZEPINE
 CARISOPRODOL, CARISOPRODOL
 CEFDINIR, CEFDINIR
 CEFPROZIL, CEFPROZIL
 CEFUROXIME AXETIL, CEFUROXIME AXETIL
 CEPHALEXIN, CEPHALEXIN
 CHLORDIAZEPOXIDE HYDROCHLORIDE AND CLIDINIUM BROMIDE, CHLORDIAZEPOXIDE HYDROCHLORIDE
 CHLORDIAZEPOXIDE HYDROCHLORIDE, CHLORDIAZEPOXIDE HYDROCHLORIDE
 CHLORHEXIDINE GLUCONATE, CHLORHEXIDINE GLUCONATE
 CHLORPROMAZINE HYDROCHLORIDE, CHLORPROMAZINE HYDROCHLORIDE
 CHLORTHALIDONE, CHLORTHALIDONE
 CICLOPIROX, CICLOPIROX
 CILOSTAZOL, CILOSTAZOL
 CINACALCET HYDROCHLORIDE, CINACALCET HYDROCHLORIDE
 CLINDAMYCIN PALMITATE HYDROCHLORIDE, CLINDAMYCIN PALMITATE HYDROCHLORIDE
 CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
 CLOMIPRAMINE HYDROCHLORIDE, CLOMIPRAMINE HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** C ****

* CHARTWELL RX SCIENCES LLC
 CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
 CYCLOBENZAPRINE HYDROCHLORIDE, CYCLOBENZAPRINE HYDROCHLORIDE
 CYPROHEPTADINE HYDROCHLORIDE, CYPROHEPTADINE HYDROCHLORIDE
 DEFERASIROX, DEFERASIROX
 DESIPRAMINE HYDROCHLORIDE, DESIPRAMINE HYDROCHLORIDE
 DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
 DIPHENOXYLATE HYDROCHLORIDE AND ATROPINE SULFATE, ATROPINE SULFATE
 DIVALPROEX SODIUM, DIVALPROEX SODIUM
 DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
 DOXEPIN HYDROCHLORIDE, DOXEPIN HYDROCHLORIDE
 DUVOID, BETHANECHOL CHLORIDE
 ELETRIPATAN HYDROBROMIDE, ELETRIPATAN HYDROBROMIDE
 EMTRICITABINE AND TENOFOVIR DISOPROXIL FUMARATE, EMTRICITABINE
 EPLERENONE, EPLERENONE
 ERGOCALCIFEROL, ERGOCALCIFEROL
 ERLOTINIB HYDROCHLORIDE, ERLOTINIB HYDROCHLORIDE
 ESCITALOPRAM OXALATE, ESCITALOPRAM OXALATE
 ETHACRYNIC ACID, ETHACRYNIC ACID
 FAMOTIDINE, FAMOTIDINE
 FENOFIBRATE, FENOFIBRATE
 FENOFIBRIC ACID, CHOLINE FENOFIBRATE
 FLUOCINOLONE ACETONIDE, FLUOCINOLONE ACETONIDE
 FLUOCINONIDE, FLUOCINONIDE
 FLUPHENAZINE HYDROCHLORIDE, FLUPHENAZINE HYDROCHLORIDE
 FLURAZEPAM HYDROCHLORIDE, FLURAZEPAM HYDROCHLORIDE
 FLUTICASONE PROPIONATE, FLUTICASONE PROPIONATE
 FOSINOPRIL SODIUM, FOSINOPRIL SODIUM
 FULVICIN P/G 165, GRISEOFULVIN, ULTRAMICROSIZE
 FULVICIN P/G 330, GRISEOFULVIN, ULTRAMICROSIZE
 FULVICIN P/G, GRISEOFULVIN, ULTRAMICROSIZE
 FULVICIN-U/F, GRISEOFULVIN, MICROSIZE
 GENERLAC, LACTULOSE
 GLYCOPYRROLATE, GLYCOPYRROLATE
 GRISEOFULVIN, GRISEOFULVIN, MICROSIZE
 HALCINONIDE, HALCINONIDE
 HALOPERIDOL, HALOPERIDOL
 HYDROXYCHLOROQUINE SULFATE, HYDROXYCHLOROQUINE SULFATE
 HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE
 IMATINIB MESYLATE, IMATINIB MESYLATE
 INDOMETHACIN, INDOMETHACIN
 KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
 LACOSAMIDE, LACOSAMIDE
 LACTULOSE, LACTULOSE
 LEVETIRACETAM, LEVETIRACETAM
 LINEZOLID, LINEZOLID
 LISINOPRIL, LISINOPRIL
 LOCHOLEST LIGHT, CHOLESTYRAMINE
 LOCHOLEST, CHOLESTYRAMINE
 LOSARTAN POTASSIUM AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 LOSARTAN POTASSIUM, LOSARTAN POTASSIUM
 LOVASTATIN, LOVASTATIN
 MECLIZINE HYDROCHLORIDE, MECLIZINE HYDROCHLORIDE
 METHAZOLAMIDE, METHAZOLAMIDE
 METHIMAZOLE, METHIMAZOLE
 METHYLERGONOVINE MALEATE, METHYLERGONOVINE MALEATE
 METRONIDAZOLE, METRONIDAZOLE
 MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE
 MIRTAZAPINE, MIRTAZAPINE
 MOEXIPRIL HYDROCHLORIDE, MOEXIPRIL HYDROCHLORIDE
 MONODOX, DOXYCYCLINE
 MOXIFLOXACIN HYDROCHLORIDE, MOXIFLOXACIN HYDROCHLORIDE
 NABUMETONE, NABUMETONE
 NADOLOL, NADOLOL
 NIACIN, NIACIN

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** C ****

* CHARTWELL RX SCIENCES LLC
 OFLOXACIN, OFLOXACIN
 OLMESARTAN MEDOXOMIL, OLMESARTAN MEDOXOMIL
 OXCARBAZEPINE, OXCARBAZEPINE
 OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE
 PALONOSETRON HYDROCHLORIDE, PALONOSETRON HYDROCHLORIDE
 PAROXETINE HYDROCHLORIDE, PAROXETINE HYDROCHLORIDE
 PENICILLIN V POTASSIUM, PENICILLIN V POTASSIUM
 PIOGLITAZONE HYDROCHLORIDE AND GLIMEPIRIDE, GLIMEPIRIDE
 PIOGLITAZONE HYDROCHLORIDE, PIOGLITAZONE HYDROCHLORIDE
 PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE
 PRAVASTATIN SODIUM, PRAVASTATIN SODIUM
 PREDNISOLONE SODIUM PHOSPHATE, PREDNISOLONE SODIUM PHOSPHATE
 PREDNISOLONE, PREDNISOLONE
 PREGABALIN, PREGABALIN
 PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
 QUINAPRIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 QUINAPRIL HYDROCHLORIDE, QUINAPRIL HYDROCHLORIDE
 RABEPRAZOLE SODIUM, RABEPRAZOLE SODIUM
 RANOLAZINE, RANOLAZINE
 RASAGILINE MESYLATE, RASAGILINE MESYLATE
 REPAGLINIDE, REPAGLINIDE
 RIVASTIGMINE TARTRATE, RIVASTIGMINE TARTRATE
 RIZATRIPTAN BENZOATE, RIZATRIPTAN BENZOATE
 ROPINIROLE HYDROCHLORIDE, ROPINIROLE HYDROCHLORIDE
 ROSUVASTATIN CALCIUM, ROSUVASTATIN CALCIUM
 RUFINAMIDE, RUFINAMIDE
 SEVELAMER CARBONATE, SEVELAMER CARBONATE
 SILDENAFIL CITRATE, SILDENAFIL CITRATE
 SILODOSIN, SILODOSIN
 SODIUM POLYSTYRENE SULFONATE, SODIUM POLYSTYRENE SULFONATE
 SOLIFENACIN SUCCINATE, SOLIFENACIN SUCCINATE
 SULFACETAMIDE SODIUM, SULFACETAMIDE SODIUM
 TADALAFIL, TADALAFIL
 TAMSULOSIN HYDROCHLORIDE, TAMSULOSIN HYDROCHLORIDE
 TAVABOROLE, TAVABOROLE
 TELMISARTAN, TELMISARTAN
 TEMAZEPAM, TEMAZEPAM
 TETRABENAZINE, TETRABENAZINE
 TINIDAZOLE, TINIDAZOLE
 TOBRAMYCIN, TOBRAMYCIN
 TRAVOPROST, TRAVOPROST
 TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE
 TRIAMTERENE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 TROSPIUM CHLORIDE, TROSPIUM CHLORIDE
 URSODIOL, URSODIOL
 VALACYCLOVIR HYDROCHLORIDE, VALACYCLOVIR HYDROCHLORIDE
 VALPROIC ACID, VALPROIC ACID
 VORICONAZOLE, VORICONAZOLE
 ZIPRASIDONE HYDROCHLORIDE, ZIPRASIDONE HYDROCHLORIDE
 ZOLEDRONIC ACID, ZOLEDRONIC ACID

CHARTWELL TETRA

* CHARTWELL TETRA LLC
 TETRACYCLINE HYDROCHLORIDE, TETRACYCLINE HYDROCHLORIDE

CHATTEM

* CHATTEM INC
 UNISOM, DOXYLAMINE SUCCINATE (OTC)

CHATTEM SANOFI

* CHATTEM INC DBA SANOFI CONSUMER HEALTHCARE
 ALLEGRA ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
 ALLEGRA HIVES, FEXOFENADINE HYDROCHLORIDE (OTC)
 ALLEGRA-D 12 HOUR ALLERGY AND CONGESTION, FEXOFENADINE HYDROCHLORIDE (OTC)
 ALLEGRA-D 24 HOUR ALLERGY AND CONGESTION, FEXOFENADINE HYDROCHLORIDE (OTC)
 CHILDREN'S ALLEGRA ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
 NASACORT ALLERGY 24 HOUR, TRIAMCINOLONE ACETONIDE (OTC)

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** C ****

- * CHATTEM INC DBA SANOFI CONSUMER HEALTHCARE
NICODERM CQ, NICOTINE (OTC)
XYZAL ALLERGY 24HR, LEVOCETIRIZINE DIHYDROCHLORIDE (OTC)

CHEMI SPA

- * CHEMI SPA
DECITABINE, DECITABINE
IBANDRONATE SODIUM, IBANDRONATE SODIUM
TEMOZOLOMIDE, TEMOZOLOMIDE

CHEMISCH FBRK KRSSLR

- * CHEMISCHE FABRIK KREUSSLER & CO. GMBH
ASCLERA, POLIDOCANOL

CHEMISTRY HLTH

- * CHEMISTRY AND HEALTH FZ LLC
CLOMIPRAMINE HYDROCHLORIDE, CLOMIPRAMINE HYDROCHLORIDE
METOLAZONE, METOLAZONE

CHEMO RESEARCH SL

- * CHEMO RESEARCH SL
BENZNIDAZOLE, BENZNIDAZOLE
ESTRADIOL, ESTRADIOL
NUVESSA, METRONIDAZOLE

CHEMOCENTRYX

- * CHEMOCENTRYX INC
TAVNEOS, AVACOPAN

CHENGDU

- * CHENGDU SUNCADIA MEDICINE CO LTD
TACROLIMUS, TACROLIMUS

CHENGDU SHUODE

- * CHENGDU SHUODE PHARMACEUTICAL CO LTD
NALMEFENE HYDROCHLORIDE, NALMEFENE HYDROCHLORIDE
NICARDIPINE HYDROCHLORIDE, NICARDIPINE HYDROCHLORIDE

CHEPLAPHARM

- * CHEPLAPHARM ARZNEIMITTEL GMBH
ETOPOPHOS PRESERVATIVE FREE, ETOPOSIDE PHOSPHATE
HYDREA, HYDROXYUREA
KLONOPIN, CLONAZEPAM
PULMICORT FLEXHALER, BUDESONIDE
VALCYTE, VALGANCICLOVIR HYDROCHLORIDE
XELODA, CAPECITABINE
XENICAL, ORLISTAT
- * CHEPLAPHARM REGISTRATION GMBH
ZYPREXA RELPREVV, OLANZAPINE PAMOATE
ZYPREXA ZYDIS, OLANZAPINE
ZYPREXA, OLANZAPINE

CHIA TAI TIANQING

- * CHIA TAI TIANQING PHARMACEUTICAL GROUP CO LTD
FOSAPREPITANT DIMEGLUMINE, FOSAPREPITANT DIMEGLUMINE
FULVESTRANT, FULVESTRANT

CHIESI

- * CHIESI FARMACEUTICI SPA
FILSUVEZ, BIRCH TRITERPENES
JUXTAPID, LOMITAPIDE MESYLATE
MYCAPSSA, OCTREOTIDE ACETATE
- * CHIESI USA INC
BETHKIS, TOBRAMYCIN
CARDENE IN 0.83% SODIUM CHLORIDE IN PLASTIC CONTAINER, NICARDIPINE HYDROCHLORIDE
CARDENE IN 0.86% SODIUM CHLORIDE IN PLASTIC CONTAINER, NICARDIPINE HYDROCHLORIDE
CARDENE IN 4.8% DEXTROSE IN PLASTIC CONTAINER, NICARDIPINE HYDROCHLORIDE
CLEVIPREX, CLEVIDIPINE
FERRIPROX, DEFERIPRONE
KENGREAL, CANGRELOR
ZYFLO, ZILEUTON

CHILDRENS HOSP MI

- * CHILDRENS HOSP MICHIGAN
FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** C ******CHINA RESOURCES**

* CHINA RESOURCES SAIKE PHARMACEUTICAL CO LTD
 AMLODIPINE BESYLATE, AMLODIPINE BESYLATE

CHIRHOCLIN

* CHIRHOCLIN INC
 CHIRHOSTIM, SECRETIN SYNTHETIC HUMAN

CINTEX SVCS

* CINTEX SERVICES LLC
 DESONIDE, DESONIDE
 TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE

CIPHER

* CIPHER PHARMACEUTICALS INC
 NATROBA, SPINOSAD

CIPHER PHARMS INC

* CIPHER PHARMACEUTICALS INC
 CONZIP, TRAMADOL HYDROCHLORIDE
 LIPOFEN, FENOFIBRATE

CIPLA

* CIPLA LTD
 ABACAVIR SULFATE AND LAMIVUDINE, ABACAVIR SULFATE
 ABACAVIR SULFATE, ABACAVIR SULFATE
 ACYCLOVIR, ACYCLOVIR
 ALBUTEROL SULFATE AND IPRATROPIUM BROMIDE, ALBUTEROL SULFATE
 ALBUTEROL SULFATE, ALBUTEROL SULFATE
 ALENDRONATE SODIUM, ALENDRONATE SODIUM
 AMBRISENTAN, AMBRISENTAN
 AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
 AMOXICILLIN AND CLAVULANATE POTASSIUM, AMOXICILLIN
 ANASTROZOLE, ANASTROZOLE
 ARFORMOTEROL TARTRATE, ARFORMOTEROL TARTRATE
 AZACITIDINE, AZACITIDINE
 BEXAROTENE, BEXAROTENE
 BUDESONIDE, BUDESONIDE
 CALCITONIN-SALMON, CALCITONIN SALMON
 CELECOXIB, CELECOXIB
 CINACALCET HYDROCHLORIDE, CINACALCET HYDROCHLORIDE
 CYCLOPHOSPHAMIDE, CYCLOPHOSPHAMIDE
 DARIFENACIN HYDROBROMIDE, DARIFENACIN HYDROBROMIDE
 DARUNAVIR, DARUNAVIR
 DEFERASIROX, DEFERASIROX
 DICLOFENAC SODIUM, DICLOFENAC SODIUM (OTC)
 DIFLUPREDNATE, DIFLUPREDNATE
 DIHYDROERGOTAMINE MESYLATE, DIHYDROERGOTAMINE MESYLATE
 DIMETHYL FUMARATE, DIMETHYL FUMARATE
 EFAVIRENZ, EFAVIRENZ
 EFAVIRENZ, EMTRICITABINE, AND TENOFOVIR DISOPROXIL FUMARATE, EFAVIRENZ
 ELIGLUSTAT TARTRATE, ELIGLUSTAT TARTRATE
 EMTRICITABINE, EMTRICITABINE
 ENTECAVIR, ENTECAVIR
 ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM
 EXEMESTANE, EXEMESTANE
 FENOFIBRATE, FENOFIBRATE
 FOSFOMYCIN TROMETHAMINE, FOSFOMYCIN TROMETHAMINE
 GRISEOFULVIN, GRISEOFULVIN, MICROSIZE
 HYDROCORTISONE SODIUM SUCCINATE, HYDROCORTISONE SODIUM SUCCINATE
 ICATIBANT ACETATE, ICATIBANT ACETATE
 LAMIVUDINE AND ZIDOVUDINE, LAMIVUDINE
 LAMIVUDINE, LAMIVUDINE
 LENALIDOMIDE, LENALIDOMIDE
 LEVALBUTEROL HYDROCHLORIDE, LEVALBUTEROL HYDROCHLORIDE
 MELOXICAM, MELOXICAM
 METOPROLOL SUCCINATE, METOPROLOL SUCCINATE
 MONTELUKAST SODIUM, MONTELUKAST SODIUM
 NEVIRAPINE, NEVIRAPINE
 NICARDIPINE HYDROCHLORIDE IN 0.9% SODIUM CHLORIDE, NICARDIPINE HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** C ****

* CIPLA LTD
 PHYTONADIONE, PHYTONADIONE
 PRAVASTATIN SODIUM, PRAVASTATIN SODIUM
 RITONAVIR, RITONAVIR
 SUMATRIPTAN, SUMATRIPTAN
 TADALAFIL, TADALAFIL
 TAVABOROLE, TAVABOROLE
 TENOFOVIR DISOPROXIL FUMARATE, TENOFOVIR DISOPROXIL FUMARATE
 TESTOSTERONE CYPIONATE, TESTOSTERONE CYPIONATE
 TESTOSTERONE, TESTOSTERONE
 VASOPRESSIN, VASOPRESSIN
 ZIDOVUDINE, ZIDOVUDINE

CIPLA LTD

* CIPLA LTD
 DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
 EPIRUBICIN HYDROCHLORIDE, EPIRUBICIN HYDROCHLORIDE
 TOPIRAMATE, TOPIRAMATE
 ZIDOVUDINE, ZIDOVUDINE

CIPLA USA

* CIPLA USA INC
 ZEMDRI, PLAZOMICIN SULFATE

CISEN

* CISEN PHARMACEUTICAL CO LTD
 ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM

CLINIGEN HLTHCARE

* CLINIGEN HEALTHCARE LTD
 FOSSCAVIR, FOSSCARNET SODIUM

CLIVUNEL INC

* CLINUNEL INC
 SCENESSE, AFAMELANOTIDE

CMP DEV LLC

* CMP DEVELOPMENT LLC
 ATORVALIQ, ATORVASTATIN CALCIUM
 CAROSPIR, SPIRONOLACTONE
 NORLIQVA, AMLODIPINE BESYLATE
 TADLIQ, TADALAFIL

CMP PHARMA INC

* CMP PHARMA INC
 HYDROCORTISONE IN ABSORBASE, HYDROCORTISONE
 ISONIAZID, ISONIAZID
 SODIUM POLYSTYRENE SULFONATE, SODIUM POLYSTYRENE SULFONATE
 SPS, SODIUM POLYSTYRENE SULFONATE

COLGATE-PALMOLIVE CO

* COLGATE-PALMOLIVE CO
 PERIOGARD, CHLORHEXIDINE GLUCONATE

COLLEGIUM PHARM INC

* COLLEGIUM PHARMACEUTICAL INC
 NUCYN TA ER, TAPENTADOL HYDROCHLORIDE
 NUCYN TA, TAPENTADOL HYDROCHLORIDE
 XTAMPZA ER, OXYCODONE

COMBE

* COMBE INC
 VAGISTAT-1, TIOCONAZOLE (OTC)

COMMAVE THERAP

* COMMAVE THERAPEUTICS SA
 AZSTARYS, DEXMETHYLPHENIDATE HYDROCHLORIDE

CONBA USA

* CONBA USA INC
 ENTECAVIR, ENTECAVIR

CONCORD BIOTECH LTD

* CONCORD BIOTECH LTD
 MYCOPHENOLATE MOFETIL, MYCOPHENOLATE MOFETIL
 MYCOPHENOLIC SODIUM, MYCOPHENOLIC SODIUM
 TACROLIMUS, TACROLIMUS

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** C ******CONTRACT PHARMACAL**

* CONTRACT PHARMACAL CORP
 CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 FAMOTIDINE, FAMOTIDINE
 FAMOTIDINE, FAMOTIDINE (OTC)
 IBUPROFEN, IBUPROFEN
 NAPROXEN SODIUM, NAPROXEN SODIUM (OTC)

COOPERSURGICAL

* COOPERSURGICAL INC
 PARAGARD T 380A, COPPER

CORCEPT THERAP

* CORCEPT THERAPEUTICS INC
 KORLYM, MIFEPRISTONE

COREPHARMA

* COREPHARMA LLC
 AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
 BENAZEPRIL HYDROCHLORIDE, BENAZEPRIL HYDROCHLORIDE
 CAPTOPRIL, CAPTOPRIL
 CHLORDIAZEPOXIDE HYDROCHLORIDE AND CLIDINIUM BROMIDE, CHLORDIAZEPOXIDE HYDROCHLORIDE
 CHLORZOXAZONE, CHLORZOXAZONE
 CLORAZEPATE DIPOTASSIUM, CLORAZEPATE DIPOTASSIUM
 DEXAMETHASONE, DEXAMETHASONE
 DICYCLOMINE HYDROCHLORIDE, DICYCLOMINE HYDROCHLORIDE
 LISINOPRIL AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 LISINOPRIL, LISINOPRIL
 LOVASTATIN, LOVASTATIN
 MELOXICAM, MELOXICAM
 NAPROXEN SODIUM AND DIPHENHYDRAMINE HYDROCHLORIDE, DIPHENHYDRAMINE HYDROCHLORIDE
 PIOGLITAZONE HYDROCHLORIDE, PIOGLITAZONE HYDROCHLORIDE
 RAMIPRIL, RAMIPRIL
 SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE

CORIUM

* CORIUM LLC
 ADLARITY, DONEPEZIL HYDROCHLORIDE

CORMEDIX

* CORMEDIX INC
 DEFENCATH, HEPARIN SODIUM

COSETTE

* COSETTE PHARMACEUTICALS INC
 ALBUTEROL SULFATE, ALBUTEROL SULFATE
 AZOR, AMLODIPINE BESYLATE
 BENICAR HCT, HYDROCHLOROTHIAZIDE
 BENICAR, OLMESARTAN MEDOXOMIL
 BETA-VAL, BETAMETHASONE VALERATE
 BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
 CALCIPOTRIENE AND BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
 CALCIPOTRIENE, CALCIPOTRIENE
 CICLOPIROX, CICLOPIROX
 CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
 CLOMIPHENE CITRATE, CLOMIPHENE CITRATE
 DAPSONE, DAPSONE
 DESONIDE, DESONIDE
 EFFIENT, PRASUGREL HYDROCHLORIDE
 EVOXAC, CEVIMELINE HYDROCHLORIDE
 FLUOCINOLONE ACETONIDE, FLUOCINOLONE ACETONIDE
 FLUOCINONIDE EMULSIFIED BASE, FLUOCINONIDE
 FLUOCINONIDE, FLUOCINONIDE
 FLUTICASONE PROPIONATE, FLUTICASONE PROPIONATE
 GENTAMICIN SULFATE, GENTAMICIN SULFATE
 GRISEOFULVIN, GRISEOFULVIN, MICROSIZE
 HALOBETASOL PROPIONATE, HALOBETASOL PROPIONATE
 HYDROCORTISONE AND ACETIC ACID, ACETIC ACID, GLACIAL
 HYDROCORTISONE VALERATE, HYDROCORTISONE VALERATE
 INDOMETHACIN, INDOMETHACIN
 KETOCONAZOLE, KETOCONAZOLE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** C ****

* COSETTE PHARMACEUTICALS INC
 M-ZOLE 3 COMBINATION PACK, MICONAZOLE NITRATE (OTC)
 METRONIDAZOLE, METRONIDAZOLE
 MICONAZOLE NITRATE, MICONAZOLE NITRATE (OTC)
 MIGERGOT, CAFFEINE
 MOMETASONE FUROATE, MOMETASONE FUROATE
 NITROGLYCERIN, NITROGLYCERIN
 NYSTATIN, NYSTATIN
 PROCHLORPERAZINE, PROCHLORPERAZINE
 PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
 PROMETHEGAN, PROMETHAZINE HYDROCHLORIDE
 SUCRALFATE, SUCRALFATE
 TAZAROTENE, TAZAROTENE
 TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE
 TRIBENZOR, AMLODIPINE BESYLATE
 VYLEESI (AUTOINJECTOR), BREMELANOTIDE ACETATE
 WELCHOL, COLESEVELAM HYDROCHLORIDE

COVIS

* COVIS PHARMA GMBH
 ALTOPREV, LOVASTATIN
 ALVESCO, CICLESONIDE
 DUAKLIR PRESSAIR, ACLIDINIUM BROMIDE
 FERAHEME, FERUMOXYTOL
 LANOXIN PEDIATRIC, DIGOXIN
 LANOXIN, DIGOXIN
 OMNARIS, CICLESONIDE
 PRILOSEC, OMEPRAZOLE MAGNESIUM
 SULAR, NISOLDIPINE
 TUDORZA PRESSAIR, ACLIDINIUM BROMIDE

CPPI CV

* CP PHARMACEUTICALS INTERNATIONAL CV
 SUTENT, SUNITINIB MALATE

CREEKWOOD PHARMS

* CREEKWOOD PHARMACEUTICALS LLC
 FENOFIBRATE, FENOFIBRATE
 HYDROXYCHLOROQUINE SULFATE, HYDROXYCHLOROQUINE SULFATE
 PREGABALIN, PREGABALIN
 RIZATRIPTAN BENZOATE, RIZATRIPTAN BENZOATE
 SILODOSIN, SILODOSIN

CROSSMEDIKA SA

* CROSSMEDIKA SA
 MEXILETINE HYDROCHLORIDE, MEXILETINE HYDROCHLORIDE
 MOXIFLOXACIN HYDROCHLORIDE, MOXIFLOXACIN HYDROCHLORIDE
 TRANLYCYPROMINE SULFATE, TRANLYCYPROMINE SULFATE
 TRAZODONE HYDROCHLORIDE, TRAZODONE HYDROCHLORIDE
 VARDENAFIL HYDROCHLORIDE, VARDENAFIL HYDROCHLORIDE

CROWN LABS

* CROWN LABORATORIES INC
 ALA-CORT, HYDROCORTISONE
 TRIDERM, TRIAMCINOLONE ACETONIDE

CROWN LABS INC

* CROWN LABORATORIES INC
 NYSTATIN AND TRIAMCINOLONE ACETONIDE, NYSTATIN
 NYSTATIN, NYSTATIN

CRYSTAL

* CRYSTAL PHARMACEUTICAL SUZHOU CO LTD
 SACUBITRIL AND VALSARTAN, SACUBITRIL

CSPC OUYI

* CSPC OUYI PHARMACEUTICAL CO LTD
 AZITHROMYCIN, AZITHROMYCIN
 CARBAMAZEPINE, CARBAMAZEPINE
 CELECOXIB, CELECOXIB
 DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE
 ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM
 GABAPENTIN, GABAPENTIN

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** C ****

* CSPC OUYI PHARMACEUTICAL CO LTD
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 PALIPERIDONE, PALIPERIDONE
 PAROXETINE HYDROCHLORIDE, PAROXETINE HYDROCHLORIDE

CSPC OUYI PHARM CO

* CSPC OUYI PHARMACEUTICAL CO LTD
 TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE

CSPC-NBP PHARM

* CSPC-NBP PHARMACEUTICAL CO LTD
 BENZONATATE, BENZONATATE
 OMEGA-3-ACID ETHYL ESTERS, OMEGA-3-ACID ETHYL ESTERS

CUBIST PHARMS LLC

* CUBIST PHARMACEUTICALS LLC
 DIFICID, FIDAXOMICIN
 SIVEXTRO, TEDIZOLID PHOSPHATE
 ZERBAXA, CEFTOLOZANE SULFATE

CUMBERLAND

* CUMBERLAND PHARMACEUTICALS INC
 SANCUSO, GRANISETRON
 VAPRISOL IN 5% DEXTROSE IN PLASTIC CONTAINER, CONIVAPTAN HYDROCHLORIDE
 VIBATIV, TELAVANCIN HYDROCHLORIDE

CUMBERLAND PHARMS

* CUMBERLAND PHARMACEUTICALS INC
 ACETADOTE, ACETYLCYSTEINE
 CALDOLOR, IBUPROFEN
 LACTULOSE, LACTULOSE

CURIUM

* CURIUM US LLC
 GALLIUM CITRATE GA 67, GALLIUM CITRATE GA-67
 IOFLUPANE I-123, IOFLUPANE I-123
 OCTREOSCAN, INDIUM IN-111 PENTETREOTIDE KIT
 SODIUM IODIDE I 123, SODIUM IODIDE I-123
 TECHNISCAN MAG3, TECHNETIUM TC-99M MERTIATIDE KIT
 TECHNISCAN PYP KIT, TECHNETIUM TC-99M PYROPHOSPHATE KIT
 TECHNISCAN, TECHNETIUM TC-99M OXIDRONATE KIT
 TECHNETIUM TC 99M SESTAMIBI, TECHNETIUM TC-99M SESTAMIBI KIT
 THALLOUS CHLORIDE TL 201, THALLOUS CHLORIDE TL-201
 ULTRA-TECHNEKOW FM, TECHNETIUM TC-99M SODIUM PERTECHNETATE GENERATOR
 ULTRATAG, TECHNETIUM TC-99M RED BLOOD CELL KIT
 XENON XE 133, XENON XE-133

CURRAX

* CURRAX PHARMACEUTICALS LLC
 ONZETRA XSAIL, SUMATRIPTAN SUCCINATE
 SILENOR, DOXEPIN HYDROCHLORIDE
 TREXIMET, NAPROXEN SODIUM

CUSTOPHARM INC

* CUSTOPHARM INC
 CALCITONIN-SALMON, CALCITONIN SALMON

CYCLE

* CYCLE PHARMACEUTICALS LTD
 NITYR, NITISINONE
 TASCENSO ODT, FINGOLIMOD LAURYL SULFATE

CYCLOMEDICA

* CYCLOMEDICA AUSTRALIA PTY LTD
 TECHNIGAS KIT, TECHNETIUM TC-99M LABELED CARBON

**** D ******DAIICHI SANKYO INC**

* DAIICHI SANKYO INC
 SAVAYSA, EDOXABAN TOSYLATE
 TURALIO, PEXIDARTINIB HYDROCHLORIDE
 VANFLYTA, QUIZARTINIB DIHYDROCHLORIDE

DAITO

* DAITO PHARMACEUTICAL CO LTD

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** D ****

- * DAITO PHARMACEUTICAL CO LTD
METHOTREXATE SODIUM, METHOTREXATE SODIUM
- DANCO LABS LLC**
- * DANCO LABORATORIES LLC
MIFEPRX, MIFEPRISTONE
- DASH PHARMS NATCO**
- * DASH PHARMACEUTICALS LLC A FULLY OWNED SUB OF NATCO PHARMA LTD
ALBUTEROL SULFATE, ALBUTEROL SULFATE
- DAVA PHARMS INC**
- * DAVA PHARMACEUTICALS INC
MORPHINE SULFATE, MORPHINE SULFATE
- DAVIS AND GECK**
- * DAVIS AND GECK DIV AMERICAN CYANAMID CO
PRE-OP II, HEXACHLOROPHENE
PRE-OP, HEXACHLOROPHENE
- DAY ONE BIOPHARMS**
- * DAY ONE BIOPHARMACEUTICALS INC
OJEMDA, TOVORAFENIB
- DECATUR**
- * DECATUR MEMORIAL HOSP
AMMONIA N 13, AMMONIA N-13
CHOLINE C-11, CHOLINE C-11
FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18
- DECIPHERA PHARMS**
- * DECIPHERA PHARMACEUTICALS LLC
QINLOCK, RIPRETINIB
- DELCATH SYSTEMS INC**
- * DELCATH SYSTEMS INC
HEPZATO, MELPHALAN HYDROCHLORIDE
- DENTSPLY PHARM**
- * DENTSPLY PHARMACEUTICAL INC
CITANEST FORTE DENTAL, EPINEPHRINE BITARTRATE
ORAQIX, LIDOCAINE
- DEPROCO**
- * DEPROCO INC
LIGNOSPAN FORTE, EPINEPHRINE BITARTRATE
LIGNOSPAN STANDARD, EPINEPHRINE BITARTRATE
SCANDONEST PLAIN, MEPIVACAINE HYDROCHLORIDE
SEPTOCAINE, ARTICAINE HYDROCHLORIDE
- DERMAVANT SCI**
- * DERMAVANT SCIENCES INC
VTAMA, TAPINAROF
- DEVA HLDING**
- * DEVA HOLDING ANONIM SIRKETI
CHLORPROMAZINE HYDROCHLORIDE, CHLORPROMAZINE HYDROCHLORIDE
- DEVA HOLDING AS**
- * DEVA HOLDING AS
AMOXICILLIN AND CLAVULANATE POTASSIUM, AMOXICILLIN
CYCLOSPORINE, CYCLOSPORINE
ESOMEPRAZOLE SODIUM, ESOMEPRAZOLE SODIUM
SUCCINYLCHOLINE CHLORIDE, SUCCINYLCHOLINE CHLORIDE
TEMOZOLOMIDE, TEMOZOLOMIDE
- DEXCEL**
- * DEXCEL PHARMA TECHNOLOGIES LTD
DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
HEMADY, DEXAMETHASONE
LANSOPRAZOLE, LANSOPRAZOLE (OTC)
OMEPRAZOLE, OMEPRAZOLE (OTC)
PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
TOFACITINIB CITRATE, TOFACITINIB CITRATE
VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE
VIGABATRIN, VIGABATRIN
- DEXCEL LTD**
- * DEXCEL LTD

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** D ****

- * DEXCEL LTD
DICLOFENAC SODIUM, DICLOFENAC SODIUM
ISOSORBIDE MONONITRATE, ISOSORBIDE MONONITRATE
- DEXCEL PHARMA**
- * DEXCEL PHARMA TECHNOLOGIES LTD
OMEPRAZOLE, OMEPRAZOLE (OTC)
- DIALYSIS SUPS**
- * DIALYSIS SUPPLIES INC
NORMOCARB HF 25, MAGNESIUM CHLORIDE
NORMOCARB HF 35, MAGNESIUM CHLORIDE
- DIFGEN PHARMS**
- * DIFGEN PHARMACEUTICALS LLC
BUPRENORPHINE HYDROCHLORIDE AND NALOXONE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE
BUPRENORPHINE, BUPRENORPHINE
CLONIDINE, CLONIDINE
FENTANYL-100, FENTANYL
FENTANYL-12, FENTANYL
FENTANYL-25, FENTANYL
FENTANYL-37, FENTANYL
FENTANYL-50, FENTANYL
FENTANYL-62, FENTANYL
FENTANYL-75, FENTANYL
FENTANYL-87, FENTANYL
NICOTINE, NICOTINE (OTC)
- DORC**
- * DORC INTERNATIONAL BV
VISIONBLUE, TRYPAN BLUE
- DOUGLAS PHARMS**
- * DOUGLAS PHARMACEUTICALS AMERICA LTD
VERSACLOZ, CLOZAPINE
- DOW PHARM**
- * DOW PHARMACEUTICAL SCIENCES
ALTRENO, TRETINOIN
ATRALIN, TRETINOIN
- DR REDDYS**
- * DR REDDYS LABORATORIES INC
CYCLOPHOSPHAMIDE, CYCLOPHOSPHAMIDE
FENOFIBRATE, FENOFIBRATE
FLUPHENAZINE HYDROCHLORIDE, FLUPHENAZINE HYDROCHLORIDE
IBUPROFEN, IBUPROFEN
IBUPROFEN, IBUPROFEN (OTC)
ICOSAPENT ETHYL, ICOSAPENT ETHYL
LOPURIN, ALLOPURINOL
METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
NITROGLYCERIN, NITROGLYCERIN
PROGESTERONE, PROGESTERONE
PROPOFOL, PROPOFOL
SSD, SILVER SULFADIAZINE
SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE
- * DR REDDYS LABORATORIES LTD
ABIRATERONE ACETATE, ABIRATERONE ACETATE
ACETAMINOPHEN AND IBUPROFEN, ACETAMINOPHEN (OTC)
ALBENDAZOLE, ALBENDAZOLE
AMLODIPINE BESYLATE AND ATORVASTATIN CALCIUM, AMLODIPINE BESYLATE
ARFORMOTEROL TARTRATE, ARFORMOTEROL TARTRATE
ASPIRIN AND DIPYRIDAMOLE, ASPIRIN
ATOMOXETINE HYDROCHLORIDE, ATOMOXETINE HYDROCHLORIDE
ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM
AZACITIDINE, AZACITIDINE
BENDAMUSTINE HYDROCHLORIDE, BENDAMUSTINE HYDROCHLORIDE
BIVALIRUDIN, BIVALIRUDIN
BORTEZOMIB, BORTEZOMIB
CABAZITAXEL, CABAZITAXEL
CALCITONIN-SALMON, CALCITONIN SALMON

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** D ****

* DR REDDYS LABORATORIES LTD
 CAPECITABINE, CAPECITABINE
 CARBOPROST TROMETHAMINE, CARBOPROST TROMETHAMINE
 CARFILZOMIB, CARFILZOMIB
 CARMUSTINE, CARMUSTINE
 CHLORDIAZEPOXIDE HYDROCHLORIDE AND CLIDINIUM BROMIDE, CHLORDIAZEPOXIDE HYDROCHLORIDE
 CHLORPROMAZINE HYDROCHLORIDE, CHLORPROMAZINE HYDROCHLORIDE
 CINACALCET HYDROCHLORIDE, CINACALCET HYDROCHLORIDE
 CIPROFLOXACIN AND DEXAMETHASONE, CIPROFLOXACIN
 CLOFARABINE, CLOFARABINE
 CLOPIDOGREL BISULFATE, CLOPIDOGREL BISULFATE
 COLCHICINE, COLCHICINE
 COLESEVELAM HYDROCHLORIDE, COLESEVELAM HYDROCHLORIDE
 DAPTOMYCIN, DAPTOMYCIN
 DARUNAVIR, DARUNAVIR
 DECITABINE, DECITABINE
 DESMOPRESSIN ACETATE, DESMOPRESSIN ACETATE
 DIAZEPAM, DIAZEPAM
 DIFLUPREDNATE, DIFLUPREDNATE
 DOCETAXEL, DOCETAXEL
 DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
 DOXORUBICIN HYDROCHLORIDE (LIPOSOMAL), DOXORUBICIN HYDROCHLORIDE
 DOXYCYCLINE, DOXYCYCLINE
 EDARAVONE, EDARAVONE
 ENALAPRILAT, ENALAPRILAT
 EPHEDRINE SULFATE, EPHEDRINE SULFATE
 ESLICARBAZEPINE ACETATE, ESLICARBAZEPINE ACETATE
 ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM
 ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM (OTC)
 ESZOPICLONE, ESZOPICLONE
 FEBUXOSTAT, FEBUXOSTAT
 FESOTERODINE FUMARATE, FESOTERODINE FUMARATE
 FEXOFENADINE HYDROCHLORIDE AND PSEUDOEPHEDRINE HYDROCHLORIDE, FEXOFENADINE
 FINGOLIMOD HYDROCHLORIDE, FINGOLIMOD HYDROCHLORIDE
 FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 FOSAPREPITANT DIMEGLUMINE, FOSAPREPITANT DIMEGLUMINE
 FULVESTRANT, FULVESTRANT
 GRANISETRON HYDROCHLORIDE, GRANISETRON HYDROCHLORIDE
 GUAIFENESIN AND DEXTROMETHORPHAN HYDROBROMIDE, DEXTROMETHORPHAN HYDROBROMIDE (OTC)
 GUAIFENESIN AND PSEUDOEPHEDRINE HYDROCHLORIDE, GUAIFENESIN (OTC)
 GUAIFENESIN, GUAIFENESIN (OTC)
 IMATINIB MESYLATE, IMATINIB MESYLATE
 LACOSAMIDE, LACOSAMIDE
 LANSOPRAZOLE, LANSOPRAZOLE
 LENALIDOMIDE, LENALIDOMIDE
 LEVOCETIRIZINE DIHYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE (OTC)
 LIDOCAINE, LIDOCAINE
 LUBIPROSTONE, LUBIPROSTONE
 LURASIDONE HYDROCHLORIDE, LURASIDONE HYDROCHLORIDE
 MELPHALAN HYDROCHLORIDE, MELPHALAN HYDROCHLORIDE
 MIDOSTAURIN, MIDOSTAURIN
 MOXIFLOXACIN HYDROCHLORIDE, MOXIFLOXACIN HYDROCHLORIDE
 NALOXONE HYDROCHLORIDE, NALOXONE HYDROCHLORIDE
 NAPROXEN AND ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM
 NELARABINE, NELARABINE
 NEOSTIGMINE METHYLSULFATE, NEOSTIGMINE METHYLSULFATE
 NYSTATIN AND TRIAMCINOLONE ACETONIDE, NYSTATIN
 OMEPRAZOLE AND SODIUM BICARBONATE, OMEPRAZOLE
 OMEPRAZOLE AND SODIUM BICARBONATE, OMEPRAZOLE (OTC)
 OMEPRAZOLE, OMEPRAZOLE (OTC)
 PALONOSETRON HYDROCHLORIDE, PALONOSETRON HYDROCHLORIDE
 PAMIDRONATE DISODIUM, PAMIDRONATE DISODIUM
 PANCURONIUM BROMIDE, PANCURONIUM BROMIDE
 PARICALCITOL, PARICALCITOL
 PEMETREXED DISODIUM, PEMETREXED DISODIUM

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** D ****

* DR REDDYS LABORATORIES LTD
 PENICILLAMINE, PENICILLAMINE
 PHYTONADIONE, PHYTONADIONE
 PLERIXAFOR, PLERIXAFOR
 POSACONAZOLE, POSACONAZOLE
 PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE
 PREGABALIN, PREGABALIN
 RABEPRAZOLE SODIUM, RABEPRAZOLE SODIUM
 REGADENOSON, REGADENOSON
 SAPROPTERIN DIHYDROCHLORIDE, SAPROPTERIN DIHYDROCHLORIDE
 SEVELAMER CARBONATE, SEVELAMER CARBONATE
 SIROLIMUS, SIROLIMUS
 SODIUM NITROPRUSSIDE, SODIUM NITROPRUSSIDE
 SORAFENIB TOSYLATE, SORAFENIB TOSYLATE
 SUCCINYLCHOLINE CHLORIDE, SUCCINYLCHOLINE CHLORIDE
 SUNITINIB MALATE, SUNITINIB MALATE
 TADALAFIL, TADALAFIL
 TETRABENAZINE, TETRABENAZINE
 THIAMINE HYDROCHLORIDE, THIAMINE HYDROCHLORIDE
 THIOTEPA, THIOTEPA
 TICAGRELOR, TICAGRELOR
 TIMOLOL MALEATE, TIMOLOL MALEATE
 TOPIRAMATE, TOPIRAMATE
 TREPROSTINIL, TREPROSTINIL
 TRIENTINE HYDROCHLORIDE, TRIENTINE HYDROCHLORIDE
 VALGANCICLOVIR HYDROCHLORIDE, VALGANCICLOVIR HYDROCHLORIDE
 VALSARTAN, VALSARTAN
 VARENICLINE TARTRATE, VARENICLINE TARTRATE
 VASOPRESSIN, VASOPRESSIN
 VIGABATRIN, VIGABATRIN
 VINOURELBINE TARTRATE, VINOURELBINE TARTRATE

DR REDDYS LABS INC

* DR REDDYS LABORATORIES INC
 AMLODIPINE BESYLATE AND BENAZEPRIL HYDROCHLORIDE, AMLODIPINE BESYLATE
 FINASTERIDE, FINASTERIDE
 FLUCONAZOLE, FLUCONAZOLE
 IBUPROFEN, IBUPROFEN
 IBUPROFEN, IBUPROFEN (OTC)
 LEVOFLOXACIN, LEVOFLOXACIN
 MELOXICAM, MELOXICAM
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 NAPROXEN SODIUM AND PSEUDOEPHEDRINE HYDROCHLORIDE, NAPROXEN SODIUM (OTC)
 NAPROXEN SODIUM, NAPROXEN SODIUM (OTC)
 PRAVASTATIN SODIUM, PRAVASTATIN SODIUM
 RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
 RIVASTIGMINE TARTRATE, RIVASTIGMINE TARTRATE
 SIMVASTATIN, SIMVASTATIN
 SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
 TERBINAFINE HYDROCHLORIDE, TERBINAFINE HYDROCHLORIDE
 TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE
 ZIPRASIDONE HYDROCHLORIDE, ZIPRASIDONE HYDROCHLORIDE

DR REDDYS LABS LTD

* DR REDDYS LABORATORIES LIMITED
 LEVOCETIRIZINE DIHYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE

* DR REDDYS LABORATORIES LTD
 ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM
 CANDESARTAN CILEXETIL AND HYDROCHLOROTHIAZIDE, CANDESARTAN CILEXETIL
 CARVEDILOL, CARVEDILOL
 CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 CETIRIZINE HYDROCHLORIDE HIVES, CETIRIZINE HYDROCHLORIDE (OTC)
 CHILDREN'S FEXOFENADINE HYDROCHLORIDE ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
 CHILDREN'S FEXOFENADINE HYDROCHLORIDE HIVES, FEXOFENADINE HYDROCHLORIDE (OTC)
 CIPROFLOXACIN EXTENDED RELEASE, CIPROFLOXACIN
 CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE
 CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** D ******* DR REDDYS LABORATORIES LTD**

CLOPIDOGREL BISULFATE, CLOPIDOGREL BISULFATE
 DESLORATADINE AND PSEUDOEPHEDRINE SULFATE 24 HOUR, DESLORATADINE
 DESLORATADINE, DESLORATADINE
 DIVALPROEX SODIUM, DIVALPROEX SODIUM
 DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
 ENALAPRIL MALEATE AND HYDROCHLOROTHIAZIDE, ENALAPRIL MALEATE
 FAMOTIDINE, FAMOTIDINE
 FAMOTIDINE, FAMOTIDINE (OTC)
 FEXOFENADINE HYDROCHLORIDE ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
 FEXOFENADINE HYDROCHLORIDE AND PSEUDOEPHEDRINE HYDROCHLORIDE, FEXOFENADINE
 FEXOFENADINE HYDROCHLORIDE HIVES, FEXOFENADINE HYDROCHLORIDE (OTC)
 FEXOFENADINE HYDROCHLORIDE, FEXOFENADINE HYDROCHLORIDE
 FINASTERIDE, FINASTERIDE
 FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 FONDAPARINUX SODIUM, FONDAPARINUX SODIUM
 GALANTAMINE HYDROBROMIDE, GALANTAMINE HYDROBROMIDE
 GEMCITABINE HYDROCHLORIDE, GEMCITABINE HYDROCHLORIDE
 GLIMEPIRIDE, GLIMEPIRIDE
 GRANISETRON HYDROCHLORIDE, GRANISETRON HYDROCHLORIDE
 IBANDRONATE SODIUM, IBANDRONATE SODIUM
 IBUPROFEN AND DIPHENHYDRAMINE CITRATE, DIPHENHYDRAMINE CITRATE (OTC)
 IBUPROFEN AND PSEUDOEPHEDRINE HYDROCHLORIDE, IBUPROFEN (OTC)
 IRBESARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 LAMOTRIGINE, LAMOTRIGINE
 LANSOPRAZOLE, LANSOPRAZOLE
 LANSOPRAZOLE, LANSOPRAZOLE (OTC)
 LEVETIRACETAM, LEVETIRACETAM
 MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
 METOPROLOL SUCCINATE, METOPROLOL SUCCINATE
 MONTELUKAST SODIUM, MONTELUKAST SODIUM
 NAPROXEN SODIUM, NAPROXEN SODIUM
 NATEGLINIDE, NATEGLINIDE
 NIZATIDINE, NIZATIDINE
 OFLOXACIN, OFLOXACIN
 OLANZAPINE, OLANZAPINE
 OMEPRAZOLE MAGNESIUM, OMEPRAZOLE MAGNESIUM (OTC)
 OMEPRAZOLE, OMEPRAZOLE
 ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
 OXAPROZIN, OXAPROZIN
 PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
 QUETIAPINE FUMARATE, QUETIAPINE FUMARATE
 RAMIPRIL, RAMIPRIL
 RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
 RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE (OTC)
 RISPERIDONE, RISPERIDONE
 ROPINIROLE HYDROCHLORIDE, ROPINIROLE HYDROCHLORIDE
 TACROLIMUS, TACROLIMUS
 TOPOTECAN HYDROCHLORIDE, TOPOTECAN HYDROCHLORIDE
 VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE
 ZAFIRLUKAST, ZAFIRLUKAST
 ZENATANE, ISOTRETINOIN
 ZOLEDRONIC ACID, ZOLEDRONIC ACID

DR REDDYS LABS SA*** DR REDDYS LABORATORIES SA**

AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE
 BIORPHEN, PHENYLEPHRINE HYDROCHLORIDE
 BRIMONIDINE TARTRATE, BRIMONIDINE TARTRATE (OTC)
 BROMPHENIRAMINE MALEATE, PSEUDOEPHEDRINE HYDROCHLORIDE AND DEXTROMETHORPHAN
 BUDESONIDE, BUDESONIDE
 BUPRENORPHINE HYDROCHLORIDE AND NALOXONE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE
 BUTALBITAL AND ACETAMINOPHEN, ACETAMINOPHEN
 BUTALBITAL, ACETAMINOPHEN AND CAFFEINE, ACETAMINOPHEN
 CAMILA, NORETHINDRONE
 CARBIDOPA AND LEVODOPA, CARBIDOPA

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** D ****

* DR REDDYS LABORATORIES SA
 CHLORZOXAZONE, CHLORZOXAZONE
 CLARITHROMYCIN, CLARITHROMYCIN
 CLONIDINE, CLONIDINE
 CYCLOSPORINE, CYCLOSPORINE
 DIAZEPAM, DIAZEPAM
 DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
 DISOPYRAMIDE PHOSPHATE, DISOPYRAMIDE PHOSPHATE
 DOFETILIDE, DOFETILIDE
 DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
 DOXYCYCLINE, DOXYCYCLINE
 DROSPIRENONE AND ETHINYL ESTRADIOL, DROSPIRENONE
 ERRIN, NORETHINDRONE
 ERYC, ERYTHROMYCIN
 ESTAZOLAM, ESTAZOLAM
 EZETIMIBE AND SIMVASTATIN, EZETIMIBE
 FENOFIBRATE (MICRONIZED), FENOFIBRATE
 FLUOROURACIL, FLUOROURACIL
 HABITROL, NICOTINE (OTC)
 HALOETTE, ETHINYL ESTRADIOL
 LEVONORGESTREL AND ETHINYL ESTRADIOL AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
 LEVONORGESTREL AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
 LEVORA 0.15/30-28, ETHINYL ESTRADIOL
 LIOTHYRONINE SODIUM, LIOTHYRONINE SODIUM
 LOW-OGESTREL-28, ETHINYL ESTRADIOL
 MERCAPTOPYRIMIDINE, MERCAPTOPYRIMIDINE
 METFORMIN HYDROCHLORIDE; SAXAGLIPTIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 METHAMPHETAMINE HYDROCHLORIDE, METHAMPHETAMINE HYDROCHLORIDE
 METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
 NICOTINE POLACRILEX, NICOTINE POLACRILEX (OTC)
 NORTRIPTYLINE HYDROCHLORIDE, NORTRIPTYLINE HYDROCHLORIDE
 NYSTATIN, NYSTATIN
 RAMELTEON, RAMELTEON
 REZIPRES, EPHEDRINE HYDROCHLORIDE
 TAMOXIFEN CITRATE, TAMOXIFEN CITRATE
 TOBRAMYCIN, TOBRAMYCIN
 TRI-NORINYL 28-DAY, ETHINYL ESTRADIOL
 TRIMETHOPRIM, TRIMETHOPRIM
 TRIVORA-28, ETHINYL ESTRADIOL

DUCHESNAY

* DUCHESNAY INC
 BONJESTA, DOXYLAMINE SUCCINATE
 DICLEGIS, DOXYLAMINE SUCCINATE
 OSPHENA, OSPHEMIFENE

DURAMED PHARMS BARR

* DURAMED PHARMACEUTICALS INC SUB BARR LABORATORIES INC
 AVIANE-28, ETHINYL ESTRADIOL
 CRYSELLE, ETHINYL ESTRADIOL
 DESOGESTREL AND ETHINYL ESTRADIOL, DESOGESTREL
 ENPRESSE-28, ETHINYL ESTRADIOL
 VELIVET, DESOGESTREL

DUTCH OPHTHALMIC

* DUTCH OPHTHALMIC RESEARCH CENTER INTERNATIONAL BV
 TISSUEBLUE, BRILLIANT BLUE G

REDDYS

* DOCTOR REDDYS LABORATORIES LTD
 DESLORATADINE, DESLORATADINE
 DIVALPROEX SODIUM, DIVALPROEX SODIUM
 METOPROLOL SUCCINATE, METOPROLOL SUCCINATE

**** E ******E5 PHARMA INC**

* E5 PHARMA INC
 DIAZOXIDE, DIAZOXIDE

EAGLE PHARMS

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** E ******* EAGLE PHARMACEUTICALS INC**

BELRAPZO, BENDAMUSTINE HYDROCHLORIDE
 BENDEKA, BENDAMUSTINE HYDROCHLORIDE
 PEMFEXY, PEMETREXED
 RYANODEX, DANTROLENE SODIUM
 VASOPRESSIN, VASOPRESSIN

ECI PHARMS LLC*** ECI PHARMACEUTICALS LLC**

TRIENTINE HYDROCHLORIDE, TRIENTINE HYDROCHLORIDE

ECOLAB*** ECOLAB INC**

CHG SCRUB, CHLORHEXIDINE GLUCONATE (OTC)
 CIDA-STAT, CHLORHEXIDINE GLUCONATE (OTC)

EDENBRIDGE PHARMS*** EDENBRIDGE PHARMACEUTICALS LLC**

ALBENDAZOLE, ALBENDAZOLE
 BISOPROLOL FUMARATE AND HYDROCHLOROTHIAZIDE, BISOPROLOL FUMARATE
 CARBIDOPA, CARBIDOPA
 COLESTIPOL HYDROCHLORIDE, COLESTIPOL HYDROCHLORIDE
 IVERMECTIN, IVERMECTIN
 LOPERAMIDE HYDROCHLORIDE, LOPERAMIDE HYDROCHLORIDE
 PREDNISOLONE SODIUM PHOSPHATE, PREDNISOLONE SODIUM PHOSPHATE
 TINIDAZOLE, TINIDAZOLE
 YARGESA, MIGLUSTAT

EIRGEN*** EIRGEN PHARMA LTD**

RAYALDEE, CALCIFEDIOL

EISAI INC*** EISAI INC**

ARICEPT, DONEPEZIL HYDROCHLORIDE
 BANZEL, RUFINAMIDE
 DAYVIGO, LEMBOREXANT
 HALAVEN, ERIBULIN MESYLATE
 LENVIMA, LENVATINIB MESYLATE

ELI LILLY AND CO*** ELI LILLY AND CO**

MOUNJARO (AUTOINJECTOR), TIRZEPATIDE
 MOUNJARO, TIRZEPATIDE
 OLUMIANT, BARICITINIB
 PROZAC, FLUOXETINE HYDROCHLORIDE
 REYVOW, LASMIDITAN SUCCINATE
 VERZENIO, ABEMACICLIB
 ZEPBOUND (AUTOINJECTOR), TIRZEPATIDE
 ZEPBOUND, TIRZEPATIDE

ELI LILLY CO*** ELI LILLY CO**

ADCIRCA, TADALAFIL

ELITE LABS*** ELITE LABORATORIES INC**

NALTREXONE HYDROCHLORIDE, NALTREXONE HYDROCHLORIDE
 PHENTERMINE HYDROCHLORIDE, PHENTERMINE HYDROCHLORIDE

ELITE LABS INC*** ELITE LABORATORIES INC**

ACETAMINOPHEN AND CODEINE PHOSPHATE, ACETAMINOPHEN
 DANTROLENE SODIUM, DANTROLENE SODIUM
 DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE
 HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
 ISRADIPINE, ISRADIPINE
 LISDEXAMFETAMINE DIMESYLATE, LISDEXAMFETAMINE DIMESYLATE
 LOXAPINE SUCCINATE, LOXAPINE SUCCINATE
 METHADONE HYDROCHLORIDE, METHADONE HYDROCHLORIDE
 METHOTREXATE SODIUM, METHOTREXATE SODIUM
 OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN
 PHENDIMETRAZINE TARTRATE, PHENDIMETRAZINE TARTRATE
 PHENTERMINE HYDROCHLORIDE, PHENTERMINE HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** E ****

* ELITE LABORATORIES INC
TRIMIPRAMINE MALEATE, TRIMIPRAMINE MALEATE

ELITE PHARM SOLUTION

* ELITE PHARMACEUTICAL SOLUTION INC
NIFEDIPINE, NIFEDIPINE

ELYSIUM

* ELYSIUM PHARMACEUTICALS LTD
AMANTADINE HYDROCHLORIDE, AMANTADINE HYDROCHLORIDE
CYPROHEPTADINE HYDROCHLORIDE, CYPROHEPTADINE HYDROCHLORIDE
POLYETHYLENE GLYCOL 3350, POLYETHYLENE GLYCOL 3350 (OTC)
TOLTERODINE TARTRATE, TOLTERODINE TARTRATE

EMD SERONO INC

* EMD SERONO INC
CETROTIDE, CETRORELIX ACETATE
MAVENCLAD, CLADRIBINE
TEPMETKO, TEPOTINIB HYDROCHLORIDE

EMED MEDCL

* EMED MEDICAL CO LLC
TERBINAFINE HYDROCHLORIDE, TERBINAFINE HYDROCHLORIDE

EMERGENT

* EMERGENT OPERATIONS IRELAND LTD
NARCAN, NALOXONE HYDROCHLORIDE (OTC)

EMERGENT BIODEFENSE

* EMERGENT BIODEFENSE OPERATIONS LANSING INC
TEMBEXA, BRINCIDOFVIR
* EMERGENT BIODEFENSE OPERATIONS LANSING LLC
TEMBEXA, BRINCIDOFVIR

EMMAUS MEDCL

* EMMAUS MEDICAL INC
ENDARI, L-GLUTAMINE

ENCUBE

* ENCUBE ETHICALS PRIVATE LTD
ADAPALENE AND BENZOYL PEROXIDE, ADAPALENE
AZELAIC ACID, AZELAIC ACID
CICLOPIROX, CICLOPIROX
CLINDAMYCIN PHOSPHATE AND BENZOYL PEROXIDE, BENZOYL PEROXIDE
CLINDAMYCIN PHOSPHATE AND TRETINOIN, CLINDAMYCIN PHOSPHATE
CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
DAPSONE, DAPSONE
DICLOFENAC SODIUM, DICLOFENAC SODIUM (OTC)
FLUOROURACIL, FLUOROURACIL
HALCINONIDE, HALCINONIDE
KETOCONAZOLE, KETOCONAZOLE
LIDOCAINE AND PRILOCAINE, LIDOCAINE
MESALAMINE, MESALAMINE
METRONIDAZOLE, METRONIDAZOLE
MOMETASONE FUROATE, MOMETASONE FUROATE
MUPIROCIN, MUPIROCIN CALCIUM
NITROGLYCERIN, NITROGLYCERIN
TACROLIMUS, TACROLIMUS
TAVABOROLE, TAVABOROLE
TESTOSTERONE, TESTOSTERONE
TRETINOIN MICROSPHERE, TRETINOIN
TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE

ENCUBE ETHICALS

* ENCUBE ETHICALS PVT LTD
ADAPALENE, ADAPALENE
CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
DESONIDE, DESONIDE
FLUOCINOLONE ACETONIDE, FLUOCINOLONE ACETONIDE
FLUOCINONIDE, FLUOCINONIDE
HYDROCORTISONE VALERATE, HYDROCORTISONE VALERATE
PERMETHRIN, PERMETHRIN

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** E ******ENDO OPERATIONS**

* ENDO OPERATIONS LTD
 ADRENALIN, EPINEPHRINE
 ALISKIREN HEMIFUMARATE, ALISKIREN HEMIFUMARATE
 ALLOPURINOL, ALLOPURINOL
 ALPRAZOLAM, ALPRAZOLAM
 ALVIMOPAN, ALVIMOPAN
 AMILORIDE HYDROCHLORIDE, AMILORIDE HYDROCHLORIDE
 ARGATROBAN, ARGATROBAN
 ASPIRIN AND DIPYRIDAMOLE, ASPIRIN
 AVEED, TESTOSTERONE UNDECANOATE
 BACLOFEN, BACLOFEN
 BENZTROPINE MESYLATE, BENZTROPINE MESYLATE
 BISMUTH SUBCITRATE POTASSIUM, METRONIDAZOLE AND TETRACYCLINE HYDROCHLORIDE, BISMUTH
 BREVITAL SODIUM, METHOHEXITAL SODIUM
 BUPRENORPHINE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE
 CALCITONIN-SALMON, CALCITONIN SALMON
 CHLORZOXAZONE, CHLORZOXAZONE
 CHOLESTYRAMINE LIGHT, CHOLESTYRAMINE
 CHOLESTYRAMINE, CHOLESTYRAMINE
 CLONAZEPAM, CLONAZEPAM
 COLCHICINE, COLCHICINE
 COLY-MYCIN M, COLISTIMETHATE SODIUM
 COLY-MYCIN S, COLISTIN SULFATE
 CORPHEDRA, EPHEDRINE SULFATE
 DANTRIUM, DANTROLENE SODIUM
 DELESTROGEN, ESTRADIOL VALERATE
 DEXLANSOPRAZOLE, DEXLANSOPRAZOLE
 DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE
 DEXMETHYLPHENIDATE HYDROCHLORIDE, DEXMETHYLPHENIDATE HYDROCHLORIDE
 DICLOFENAC POTASSIUM, DICLOFENAC POTASSIUM
 DICYCLOMINE HYDROCHLORIDE, DICYCLOMINE HYDROCHLORIDE
 DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
 DOXEPIN HYDROCHLORIDE, DOXEPIN HYDROCHLORIDE
 DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
 DOXYLAMINE SUCCINATE AND PYRIDOXINE HYDROCHLORIDE, DOXYLAMINE SUCCINATE
 DUTASTERIDE AND TAMSULOSIN HYDROCHLORIDE, DUTASTERIDE
 EDEX, ALPROSTADIL
 EPHEDRINE SULFATE, EPHEDRINE SULFATE
 ETHACRYNATE SODIUM, ETHACRYNATE SODIUM
 EVEROLIMUS, EVEROLIMUS
 FAMOTIDINE; IBUPROFEN, FAMOTIDINE
 FELODIPINE, FELODIPINE
 FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 FLUPHENAZINE DECANOATE, FLUPHENAZINE DECANOATE
 FLUVOXAMINE MALEATE, FLUVOXAMINE MALEATE
 FROVA, FROVATRIPTAN SUCCINATE
 GANCICLOVIR SODIUM, GANCICLOVIR SODIUM
 GILDAGIA, ETHINYL ESTRADIOL
 GILDESS 24 FE, ETHINYL ESTRADIOL
 GLYCOPYRROLATE, GLYCOPYRROLATE
 HYDROXYUREA, HYDROXYUREA
 IBUPROFEN, IBUPROFEN (OTC)
 ISOSORBIDE DINITRATE, ISOSORBIDE DINITRATE
 KETALAR, KETAMINE HYDROCHLORIDE
 LABETALOL HYDROCHLORIDE, LABETALOL HYDROCHLORIDE
 LAMOTRIGINE, LAMOTRIGINE
 MEGACE ES, MEGESTROL ACETATE
 METHYLPREDNISOLONE ACETATE, METHYLPREDNISOLONE ACETATE
 METHYLPREDNISOLONE, METHYLPREDNISOLONE
 MICAFUNGIN, MICAFUNGIN SODIUM
 MIDODRINE HYDROCHLORIDE, MIDODRINE HYDROCHLORIDE
 MINOXIDIL, MINOXIDIL
 MYCOPHENOLATE MOFETIL HYDROCHLORIDE, MYCOPHENOLATE MOFETIL HYDROCHLORIDE
 MYZILRA, ETHINYL ESTRADIOL

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** E ****

* ENDO OPERATIONS LTD
 NALOXONE HYDROCHLORIDE, NALOXONE HYDROCHLORIDE
 NEOSTIGMINE METHYLSULFATE, NEOSTIGMINE METHYLSULFATE
 OLANZAPINE AND FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 OLMESARTAN MEDOXOMIL, AMLODIPINE AND HYDROCHLOROTHIAZIDE, AMLODIPINE BESYLATE
 PENICILLAMINE, PENICILLAMINE
 PERCOCET, ACETAMINOPHEN
 PERCODAN, ASPIRIN
 PERPHENAZINE, PERPHENAZINE
 PHENOXYBENZAMINE HYDROCHLORIDE, PHENOXYBENZAMINE HYDROCHLORIDE
 PHENYLEPHRINE HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE
 PIMOZIDE, PIMOZIDE
 PITOCIN, OXYTOCIN
 POSACONAZOLE, POSACONAZOLE
 PRAZIQUANTEL, PRAZIQUANTEL
 PREDNISON, PREDNISON
 PREVIFEM, ETHINYL ESTRADIOL
 PROPRANOLOL HYDROCHLORIDE, PROPRANOLOL HYDROCHLORIDE
 PROPYLTHIOURACIL, PROPYLTHIOURACIL
 RISPERIDONE, RISPERIDONE
 SAPROPTERIN DIHYDROCHLORIDE, SAPROPTERIN DIHYDROCHLORIDE
 SODIUM PHENYLBUTYRATE, SODIUM PHENYLBUTYRATE
 SUPPRELIN LA, HISTRELIN ACETATE
 TESTIM, TESTOSTERONE
 TESTOPEL, TESTOSTERONE
 THEO-24, THEOPHYLLINE
 TIGAN, TRIMETHOBENZAMIDE HYDROCHLORIDE
 TIOPRONIN, TIOPRONIN
 TOLVAPTAN, TOLVAPTAN
 TOPIRAMATE, TOPIRAMATE
 TREPROSTINIL, TREPROSTINIL
 TRETINOIN, TRETINOIN
 TRIENTINE HYDROCHLORIDE, TRIENTINE HYDROCHLORIDE
 URSODIOL, URSODIOL
 VALSTAR PRESERVATIVE FREE, VALRUBICIN
 VARENICLINE TARTRATE, VARENICLINE TARTRATE
 VASOSTRICT, VASOPRESSIN
 VIGABATRIN, VIGABATRIN
 ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE

ENTASIS THERAP

* ENTASIS THERAPEUTICS INC
 XACDURO (COPACKAGED), DURLOBACTAM SODIUM

EPIC PHARMA

* EPIC PHARMA LLC
 BETAXOLOL HYDROCHLORIDE, BETAXOLOL HYDROCHLORIDE
 CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE
 FLAVOXATE HYDROCHLORIDE, FLAVOXATE HYDROCHLORIDE
 SULINDAC, SULINDAC
 TRANDOLAPRIL, TRANDOLAPRIL
 URSODIOL, URSODIOL

EPIC PHARMA LLC

* EPIC PHARMA LLC
 ACETAZOLAMIDE, ACETAZOLAMIDE
 AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
 AMPHETAMINE SULFATE, AMPHETAMINE SULFATE
 AZELASTINE HYDROCHLORIDE, AZELASTINE HYDROCHLORIDE
 AZITHROMYCIN, AZITHROMYCIN
 BENZPHETAMINE HYDROCHLORIDE, BENZPHETAMINE HYDROCHLORIDE
 BENZTROPINE MESYLATE, BENZTROPINE MESYLATE
 BISOPROLOL FUMARATE AND HYDROCHLOROTHIAZIDE, BISOPROLOL FUMARATE
 BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
 BUSPIRONE HYDROCHLORIDE, BUSPIRONE HYDROCHLORIDE
 CARBAMAZEPINE, CARBAMAZEPINE
 CARBOPLATIN, CARBOPLATIN
 CEFTRIAZONE, CEFTRIAZONE SODIUM

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** E ****

* EPIC PHARMA LLC
 CHOLESTYRAMINE LIGHT, CHOLESTYRAMINE
 CHOLESTYRAMINE, CHOLESTYRAMINE
 CILOSTAZOL, CILOSTAZOL
 CLINDAMYCIN HYDROCHLORIDE, CLINDAMYCIN HYDROCHLORIDE
 CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
 CYCLOPHOSPHAMIDE, CYCLOPHOSPHAMIDE
 DEMECLOCYCLINE HYDROCHLORIDE, DEMECLOCYCLINE HYDROCHLORIDE
 DESOXIMETASONE, DESOXIMETASONE
 DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE
 DICLOFENAC SODIUM, DICLOFENAC SODIUM
 DIFLUPREDNATE, DIFLUPREDNATE
 DORZOLAMIDE HYDROCHLORIDE AND TIMOLOL MALEATE, DORZOLAMIDE HYDROCHLORIDE
 DOXEPIN HYDROCHLORIDE, DOXEPIN HYDROCHLORIDE
 DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
 EPIRUBICIN HYDROCHLORIDE, EPIRUBICIN HYDROCHLORIDE
 ERYTHROMYCIN, ERYTHROMYCIN
 ESTRADIOL, ESTRADIOL
 ETHAMBUTOL HYDROCHLORIDE, ETHAMBUTOL HYDROCHLORIDE
 ETHOSUXIMIDE, ETHOSUXIMIDE
 FAMOTIDINE, FAMOTIDINE
 FUROSEMIDE, FUROSEMIDE
 GEMFIBROZIL, GEMFIBROZIL
 GLIPIZIDE AND METFORMIN HYDROCHLORIDE, GLIPIZIDE
 GLYBURIDE, GLYBURIDE
 GUANFACINE HYDROCHLORIDE, GUANFACINE HYDROCHLORIDE
 HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
 HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE
 IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE
 LABETALOL HYDROCHLORIDE, LABETALOL HYDROCHLORIDE
 LAMIVUDINE AND ZIDOVUDINE, LAMIVUDINE
 LEUCOVORIN CALCIUM, LEUCOVORIN CALCIUM
 LOVASTATIN, LOVASTATIN
 MECLIZINE HYDROCHLORIDE, MECLIZINE HYDROCHLORIDE
 MEPERIDINE HYDROCHLORIDE, MEPERIDINE HYDROCHLORIDE
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 METHADONE HYDROCHLORIDE, METHADONE HYDROCHLORIDE
 MIDAZOLAM HYDROCHLORIDE, MIDAZOLAM HYDROCHLORIDE
 MOLINDONE HYDROCHLORIDE, MOLINDONE HYDROCHLORIDE
 NICARDIPINE HYDROCHLORIDE, NICARDIPINE HYDROCHLORIDE
 NIZATIDINE, NIZATIDINE
 NYSTATIN AND TRIAMCINOLONE ACETONIDE, NYSTATIN
 NYSTATIN, NYSTATIN
 OLANZAPINE AND FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 OSELTAMIVIR PHOSPHATE, OSELTAMIVIR PHOSPHATE
 OXAZEPAM, OXAZEPAM
 OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN
 OXYCODONE AND ASPIRIN, ASPIRIN
 OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
 OXYMORPHONE HYDROCHLORIDE, OXYMORPHONE HYDROCHLORIDE
 PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
 PHENYTOIN, PHENYTOIN
 POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
 PREGABALIN, PREGABALIN
 PROTRIPTYLINE HYDROCHLORIDE, PROTRIPTYLINE HYDROCHLORIDE
 QUINIDINE SULFATE, QUINIDINE SULFATE
 RIFAMPIN, RIFAMPIN
 SODIUM POLYSTYRENE SULFONATE, SODIUM POLYSTYRENE SULFONATE
 SOTALOL HYDROCHLORIDE, SOTALOL HYDROCHLORIDE
 SULFADIAZINE, SULFADIAZINE
 TIMOLOL MALEATE, TIMOLOL MALEATE
 TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE
 TRANEXAMIC ACID, TRANEXAMIC ACID
 TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE
 TRIMETHOPRIM SULFATE AND POLYMYXIN B SULFATE, POLYMYXIN B SULFATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** E ****

* EPIC PHARMA LLC
 TROPICACYL, TROPICAMIDE
 URSODIOL, URSODIOL
 VORICONAZOLE, VORICONAZOLE
 ZOLEDRONIC ACID, ZOLEDRONIC ACID

EPIZYME INC

* EPIZYME INC
 TAZVERIK, TAZEMETOSTAT HYDROBROMIDE

ESJAY PHARMA

* ESJAY PHARMA LLC
 CLARITHROMYCIN, CLARITHROMYCIN
 DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
 DRISDOL, ERGOCALCIFEROL
 HIPREX, METHENAMINE HIPPURATE
 MELOXICAM, MELOXICAM
 METRONIDAZOLE, METRONIDAZOLE
 PARLODEL, BROMOCRIPTINE MESYLATE
 RISPERIDONE, RISPERIDONE
 ROCALTROL, CALCITRIOL
 SOLIFENACIN SUCCINATE, SOLIFENACIN SUCCINATE
 TETRACYCLINE HYDROCHLORIDE, TETRACYCLINE HYDROCHLORIDE

ESKAYEF

* ESKAYEF PHARMACEUTICALS LTD
 PREGABALIN, PREGABALIN

ESPERION THERAPS INC

* ESPERION THERAPEUTICS INC
 NEXLETOL, BEMPEDOIC ACID
 NEXLIZET, BEMPEDOIC ACID

ESSENTIAL ISOTOPES

* ESSENTIAL ISOTOPES LLC
 SODIUM FLUORIDE F-18, SODIUM FLUORIDE F-18

ETHYPHARM

* ETHYPHARM
 BUPRENORPHINE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE
 * ETHYPHARM SA
 ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM

ETHYPHARM USA CORP

* ETHYPHARM USA CORP
 BUPRENORPHINE HYDROCHLORIDE AND NALOXONE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE

ETON

* ETON PHARMACEUTICALS INC
 ALKINDI SPRINKLE, HYDROCORTISONE
 BETAINE, BETAINE
 NITISINONE, NITISINONE

EUGIA PHARMA

* EUGIA PHARMA SPECIALITIES LTD
 ACETAMINOPHEN, ACETAMINOPHEN
 ACETYLCYSTEINE, ACETYLCYSTEINE
 ACYCLOVIR SODIUM, ACYCLOVIR SODIUM
 ADENOSINE, ADENOSINE
 AMPHOTERICIN B, AMPHOTERICIN B
 ANASTROZOLE, ANASTROZOLE
 ATRACURIUM BESYLATE PRESERVATIVE FREE, ATRACURIUM BESYLATE
 ATRACURIUM BESYLATE, ATRACURIUM BESYLATE
 AZACITIDINE, AZACITIDINE
 AZITHROMYCIN, AZITHROMYCIN
 BENDAMUSTINE HYDROCHLORIDE, BENDAMUSTINE HYDROCHLORIDE
 BIMATOPROST, BIMATOPROST
 BIVALIRUDIN, BIVALIRUDIN
 BORTEZOMIB, BORTEZOMIB
 BROMFENAC SODIUM, BROMFENAC SODIUM
 BUPIVACAINE HYDROCHLORIDE PRESERVATIVE FREE, BUPIVACAINE HYDROCHLORIDE
 BUPIVACAINE HYDROCHLORIDE, BUPIVACAINE HYDROCHLORIDE
 CAFFEINE CITRATE, CAFFEINE CITRATE
 CAPECITABINE, CAPECITABINE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** E ****

* EUGIA PHARMA SPECIALITIES LTD
 CARBOPLATIN, CARBOPLATIN
 CARBOPROST TROMETHAMINE, CARBOPROST TROMETHAMINE
 CHLORPROMAZINE HYDROCHLORIDE, CHLORPROMAZINE HYDROCHLORIDE
 CISATRACURIUM BESYLATE PRESERVATIVE FREE, CISATRACURIUM BESYLATE
 CISATRACURIUM BESYLATE, CISATRACURIUM BESYLATE
 DACTINOMYCIN, DACTINOMYCIN
 DAPTOMYCIN, DAPTOMYCIN
 DECITABINE, DECITABINE
 DEXAMETHASONE SODIUM PHOSPHATE, DEXAMETHASONE SODIUM PHOSPHATE
 DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE
 DEXRAZOXANE HYDROCHLORIDE, DEXRAZOXANE HYDROCHLORIDE
 DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
 DOCETAXEL, DOCETAXEL
 DORZOLAMIDE HYDROCHLORIDE AND TIMOLOL MALEATE, DORZOLAMIDE HYDROCHLORIDE
 DOXERCALCIFEROL, DOXERCALCIFEROL
 EPHEDRINE SULFATE, EPHEDRINE SULFATE
 EPTIFIBATIDE, EPTIFIBATIDE
 ERTAPENEM SODIUM, ERTAPENEM SODIUM
 ESMOLOL HYDROCHLORIDE, ESMOLOL HYDROCHLORIDE
 ESOMEPRAZOLE SODIUM, ESOMEPRAZOLE SODIUM
 ETOMIDATE, ETOMIDATE
 EXEMESTANE, EXEMESTANE
 FLUPHENAZINE DECANOATE, FLUPHENAZINE DECANOATE
 FONDAPARINUX SODIUM, FONDAPARINUX SODIUM
 FOSAPREPITANT DIMEGLUMINE, FOSAPREPITANT DIMEGLUMINE
 FULVESTRANT, FULVESTRANT
 FUROSEMIDE, FUROSEMIDE
 GENTAMICIN SULFATE, GENTAMICIN SULFATE
 HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
 IBANDRONATE SODIUM, IBANDRONATE SODIUM
 ICATIBANT ACETATE, ICATIBANT ACETATE
 IMATINIB MESYLATE, IMATINIB MESYLATE
 IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE
 KETAMINE HYDROCHLORIDE, KETAMINE HYDROCHLORIDE
 LENALIDOMIDE, LENALIDOMIDE
 LETROZOLE, LETROZOLE
 LEUPROLIDE ACETATE, LEUPROLIDE ACETATE
 LEVETIRACETAM IN SODIUM CHLORIDE, LEVETIRACETAM
 LEVETIRACETAM, LEVETIRACETAM
 LIDOCAINE HYDROCHLORIDE PRESERVATIVE FREE, LIDOCAINE HYDROCHLORIDE
 LIDOCAINE HYDROCHLORIDE, LIDOCAINE HYDROCHLORIDE
 LINEZOLID, LINEZOLID
 MEDROXYPROGESTERONE ACETATE, MEDROXYPROGESTERONE ACETATE
 MEROPENEM, MEROPENEM
 METHOCARBAMOL, METHOCARBAMOL
 METHOTREXATE SODIUM, METHOTREXATE SODIUM
 METHYLPREDNISOLONE ACETATE, METHYLPREDNISOLONE ACETATE
 MILRINONE LACTATE IN DEXTROSE 5%, MILRINONE LACTATE
 MITOMYCIN, MITOMYCIN
 MOXIFLOXACIN HYDROCHLORIDE, MOXIFLOXACIN HYDROCHLORIDE
 NALOXONE HYDROCHLORIDE, NALOXONE HYDROCHLORIDE
 NICARDIPINE HYDROCHLORIDE, NICARDIPINE HYDROCHLORIDE
 OLANZAPINE, OLANZAPINE
 OLOPATADINE HYDROCHLORIDE, OLOPATADINE HYDROCHLORIDE (OTC)
 PALONOSETRON HYDROCHLORIDE, PALONOSETRON HYDROCHLORIDE
 PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
 PARICALCITOL, PARICALCITOL
 PAZOPANIB HYDROCHLORIDE, PAZOPANIB HYDROCHLORIDE
 PEMETREXED DISODIUM, PEMETREXED DISODIUM
 PHENYLEPHRINE HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE
 PLERIXAFOR, PLERIXAFOR
 POLYMYXIN B SULFATE, POLYMYXIN B SULFATE
 POSACONAZOLE, POSACONAZOLE
 PROCHLORPERAZINE EDISYLATE, PROCHLORPERAZINE EDISYLATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** E ****

* EUGIA PHARMA SPECIALITIES LTD
 PROGESTERONE, PROGESTERONE
 REGADENOSON, REGADENOSON
 ROCURONIUM BROMIDE, ROCURONIUM BROMIDE
 ROPIVACAINE HYDROCHLORIDE, ROPIVACAINE HYDROCHLORIDE
 SILDENAFIL CITRATE, SILDENAFIL CITRATE
 SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
 SUNITINIB MALATE, SUNITINIB MALATE
 TAMOXIFEN CITRATE, TAMOXIFEN CITRATE
 TESTOSTERONE CYPIONATE, TESTOSTERONE CYPIONATE
 TESTOSTERONE ENANTHATE, TESTOSTERONE ENANTHATE
 THIAMINE HYDROCHLORIDE, THIAMINE HYDROCHLORIDE
 TIGECYCLINE, TIGECYCLINE
 TIMOLOL MALEATE, TIMOLOL MALEATE
 TIROFIBAN HYDROCHLORIDE, TIROFIBAN HYDROCHLORIDE
 TOBRAMYCIN SULFATE, TOBRAMYCIN SULFATE
 TRANEXAMIC ACID, TRANEXAMIC ACID
 TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE
 VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE
 VASOPRESSIN, VASOPRESSIN
 VECURONIUM BROMIDE, VECURONIUM BROMIDE
 VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE

EUGIA PHARMA SPECLTS

* EUGIA PHARMA SPECIALITIES LTD
 AMPICILLIN AND SULBACTAM, AMPICILLIN SODIUM
 AMPICILLIN SODIUM, AMPICILLIN SODIUM
 ARGATROBAN IN SODIUM CHLORIDE, ARGATROBAN
 CYCLOPHOSPHAMIDE, CYCLOPHOSPHAMIDE
 FLUOROURACIL, FLUOROURACIL
 NAFCILLIN SODIUM, NAFCILLIN SODIUM
 OXACILLIN SODIUM, OXACILLIN SODIUM
 PIPERACILLIN AND TAZOBACTAM, PIPERACILLIN SODIUM

EUROCEPT PHARMS

* EUROCEPT PHARMACEUTICALS
 PASER, AMINOSALICYLIC ACID

EUROHLTH INTL SARL

* EUROHEALTH INTERNATIONAL SARL
 AZACITIDINE, AZACITIDINE

EVEREST LIFE SCI

* EVEREST LIFE SCIENCES LLC
 DAPSONE, DAPSONE

EVOFEM INC

* EVOFEM INC
 PHEXXI, CITRIC ACID
 SOLOSEC, SECNIDAZOLE

EVOKE PHARMA INC

* EVOKE PHARMA INC
 GIMOTI, METOCLOPRAMIDE HYDROCHLORIDE

EVUS

* EVUS HEALTH SOLUTIONS LLC
 NITROMIST, NITROGLYCERIN

EXELA PHARMA

* EXELA PHARMA SCIENCES LLC
 AKOVAZ, EPHEDRINE SULFATE
 BLOXIVERZ, NEOSTIGMINE METHYLSULFATE
 CAFFEINE CITRATE, CAFFEINE CITRATE
 CUPRIC CHLORIDE, CUPRIC CHLORIDE
 ELCYS, CYSTEINE HYDROCHLORIDE
 GANZYK-RTU, GANCICLOVIR
 GLYRX-PF, GLYCOPYRROLATE
 NIPRIDE RTU IN SODIUM CHLORIDE 0.9%, SODIUM NITROPRUSSIDE
 POTASSIUM ACETATE, POTASSIUM ACETATE
 SODIUM BICARBONATE, SODIUM BICARBONATE
 TRANEXAMIC ACID, TRANEXAMIC ACID
 VAZCULEP, PHENYLEPHRINE HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** E ****

* EXELA PHARMA SCIENCES LLC
 VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE
 ZINC CHLORIDE, ZINC CHLORIDE

EXELA PHARMA SCIENCE

* EXELA PHARMA SCIENCES
 CAFFEINE CITRATE, CAFFEINE CITRATE

EXELIXIS

* EXELIXIS INC
 COMETRIQ, CABOZANTINIB S-MALATE

EXELIXIS INC

* EXELIXIS INC
 CABOMETYX, CABOZANTINIB S-MALATE

EXELTIS USA INC

* EXELTIS USA INC
 SLYND, DROSPIRENONE
 TYBLUME, ETHINYL ESTRADIOL

EXTROVIS

* EXTROVIS AG
 CALCIUM CHLORIDE 10%, CALCIUM CHLORIDE
 CARAC, FLUOROURACIL
 EFUDEX, FLUOROURACIL

EYENOVIA

* EYENOVIA INC
 CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE

EYEPOINT PHARMS

* EYEPOINT PHARMACEUTICALS INC
 DEXYCU KIT, DEXAMETHASONE

EZRA VENTURES

* EZRA VENTURES LLC
 FINGOLIMOD HYDROCHLORIDE, FINGOLIMOD HYDROCHLORIDE

LILLY

* ELI LILLY AND CO
 ALIMTA, PEMETREXED DISODIUM
 CIALIS, TADALAFIL
 CYMBALTA, DULOXETINE HYDROCHLORIDE
 EVISTA, RALOXIFENE HYDROCHLORIDE
 FORTEO, TERIPARATIDE
 STRATTERA, ATOMOXETINE HYDROCHLORIDE
 SYMBYAX, FLUOXETINE HYDROCHLORIDE

**** F ******FAMYGEN LIFE SCI**

* FAMYGEN LIFE SCIENCES INC
 RYZUMVI, PHENTOLAMINE MESYLATE

FDC LTD

* FDC LTD
 CEFIXIME, CEFIXIME
 CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE
 DORZOLAMIDE HYDROCHLORIDE AND TIMOLOL MALEATE, DORZOLAMIDE HYDROCHLORIDE
 DORZOLAMIDE HYDROCHLORIDE, DORZOLAMIDE HYDROCHLORIDE
 LATANOPROST, LATANOPROST
 OFLOXACIN, OFLOXACIN
 TIMOLOL MALEATE, TIMOLOL MALEATE

FDN CONSUMER

* FOUNDATION CONSUMER BRANDS LLC
 ALAVERT, LORATADINE (OTC)
 * FOUNDATION CONSUMER HEALTHCARE LLC
 PLAN B ONE-STEP, LEVONORGESTREL (OTC)

FEINSTEIN

* FEINSTEIN INSTITUTE MEDICAL RESEARCH
 AMMONIA N 13, AMMONIA N-13
 FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18
 FLUORODOPA F18, FLUORODOPA F-18

FENNEC PHARMS INC

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** F ****

* FENNEC PHARMACEUTICALS INC
PEDMARK, SODIUM THIOSULFATE

FERA PHARMS LLC

* FERA PHARMACEUTICALS LLC
GENTAMICIN SULFATE, GENTAMICIN SULFATE
PHOSPHOLINE IODIDE, ECHOTHIOPHATE IODIDE

FERRING

* FERRING PHARMACEUTICALS INC
ENDOMETRIN, PROGESTERONE
FIRMAGON, DEGARELIX ACETATE

FERRING PHARMS INC

* FERRING PHARMACEUTICALS INC
CERVIDIL, DINOPROSTONE
CLENPIQ, CITRIC ACID
DDAVP, DESMOPRESSIN ACETATE

FIDELITY BIOPHARMA

* FIDELITY BIOPHARMA CO USA
ONTRALFY, TIZANIDINE HYDROCHLORIDE

FLORIDA

* FLORIDA PHARMACEUTICAL PRODUCTS LLC
ABIRATERONE ACETATE, ABIRATERONE ACETATE
AMINOCAPROIC ACID, AMINOCAPROIC ACID
BRIMONIDINE TARTRATE AND TIMOLOL MALEATE, BRIMONIDINE TARTRATE

FOLDRX PHARMS

* FOLDRX PHARMACEUTICALS LLC A WHOLLY OWNED SUB OF PFIZER INC
VYNDAMAX, TAFAMIDIS
VYNDAQEL, TAFAMIDIS MEGLUMINE

FOSUN PHARMA

* FOSUN PHARMA USA INC
DICYCLOMINE HYDROCHLORIDE, DICYCLOMINE HYDROCHLORIDE

FOUGERA PHARMS

* FOUGERA PHARMACEUTICALS INC
ADAPALENE, ADAPALENE
ALCLOMETASONE DIPROPIONATE, ALCLOMETASONE DIPROPIONATE
BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
CALCIPOTRIENE, CALCIPOTRIENE
CICLOPIROX, CICLOPIROX
CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
CLOBETASOL PROPIONATE (EMOLLIENT), CLOBETASOL PROPIONATE
CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
CLOTRIMAZOLE AND BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
CLOTRIMAZOLE, CLOTRIMAZOLE
DESONIDE, DESONIDE
DESOXIMETASONE, DESOXIMETASONE
ERYTHROMYCIN, ERYTHROMYCIN
FLUOCINONIDE EMULSIFIED BASE, FLUOCINONIDE
FLUOCINONIDE, FLUOCINONIDE
FLUTICASONE PROPIONATE, FLUTICASONE PROPIONATE
HALOBETASOL PROPIONATE, HALOBETASOL PROPIONATE
HYDROCORTISONE, HYDROCORTISONE
IMIQUIMOD, IMIQUIMOD
KETOCONAZOLE, KETOCONAZOLE
LIDOCAINE AND PRILOCAINE, LIDOCAINE
METRONIDAZOLE, METRONIDAZOLE
MOMETASONE FUROATE, MOMETASONE FUROATE
MUPIROCIN, MUPIROCIN
NYSTATIN, NYSTATIN
OXISTAT, OXICONAZOLE NITRATE
PREDNICARBATE, PREDNICARBATE
SULFACETAMIDE SODIUM, SULFACETAMIDE SODIUM
TERCONAZOLE, TERCONAZOLE
TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE
TYZINE, TETRAHYDROZOLINE HYDROCHLORIDE

FOUGERA PHARMS INC

* FOUGERA PHARMACEUTICALS INC

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** F ****

* FOUGERA PHARMACEUTICALS INC
 ACYCLOVIR, ACYCLOVIR
 BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
 BETAMETHASONE VALERATE, BETAMETHASONE VALERATE
 CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
 CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
 FLUOCINOLONE ACETONIDE, FLUOCINOLONE ACETONIDE
 FLUOCINONIDE, FLUOCINONIDE
 GENTAMICIN SULFATE, GENTAMICIN SULFATE
 HYDROCORTISONE, HYDROCORTISONE
 LIDOCAINE, LIDOCAINE
 NITROGLYCERIN, NITROGLYCERIN
 NYSTATIN AND TRIAMCINOLONE ACETONIDE, NYSTATIN
 NYSTATIN, NYSTATIN
 TACROLIMUS, TACROLIMUS
 TERCONAZOLE, TERCONAZOLE

FOURRTS LABS

* FOURRTS INDIA LABORATORIES PRIVATE LTD
 TADALAFIL, TADALAFIL

FRESENIUS

* FRESENIUS KABI DEUTSCHLAND GMBH
 INTRALIPID 20%, SOYBEAN OIL
 INTRALIPID 30%, SOYBEAN OIL
 * FRESENIUS KABI IPSUM SRL
 NAFCILLIN SODIUM, NAFCILLIN SODIUM

FRESENIUS KABI

* FRESENIUS KABI ANTI INFECTIVES SRL
 PIPERACILLIN AND TAZOBACTAM, PIPERACILLIN SODIUM
 * FRESENIUS KABI AUSTRIA GMBH
 LACTULOSE, LACTULOSE

FRESENIUS KABI USA

* FRESENIUS KABI USA LLC
 ACETAMINOPHEN, ACETAMINOPHEN
 ACETYLCYSTEINE, ACETYLCYSTEINE
 ACYCLOVIR SODIUM, ACYCLOVIR SODIUM
 ADENOSINE, ADENOSINE
 AMIKACIN SULFATE, AMIKACIN SULFATE
 AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE
 ARGATROBAN, ARGATROBAN
 ARSENIC TRIOXIDE, ARSENIC TRIOXIDE
 ATROPINE SULFATE, ATROPINE SULFATE
 AZITHROMYCIN, AZITHROMYCIN
 AZTREONAM, AZTREONAM
 BACTERIOSTATIC SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, SODIUM CHLORIDE
 BENZTROPINE MESYLATE, BENZTROPINE MESYLATE
 BIVALIRUDIN, BIVALIRUDIN
 BLEOMYCIN SULFATE, BLEOMYCIN SULFATE
 BORTEZOMIB, BORTEZOMIB
 CAFFEINE CITRATE, CAFFEINE CITRATE
 CALCITONIN-SALMON, CALCITONIN SALMON
 CALCIUM GLUCONATE IN SODIUM CHLORIDE, CALCIUM GLUCONATE
 CALCIUM GLUCONATE, CALCIUM GLUCONATE
 CARBOPLATIN, CARBOPLATIN
 CARDIOPLEGIC IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 CASPOFUNGIN ACETATE, CASPOFUNGIN ACETATE
 CEFOTETAN, CEFOTETAN DISODIUM
 CHLOROTHIAZIDE SODIUM, CHLOROTHIAZIDE SODIUM
 CISATRACURIUM BESYLATE PRESERVATIVE FREE, CISATRACURIUM BESYLATE
 CISATRACURIUM BESYLATE, CISATRACURIUM BESYLATE
 CISPLATIN, CISPLATIN
 CLADRIBINE, CLADRIBINE
 CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
 CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
 COLISTIMETHATE SODIUM, COLISTIMETHATE SODIUM
 CYTARABINE, CYTARABINE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** F **

* FRESENIUS KABI USA LLC
 DACARBAZINE, DACARBAZINE
 DAPTOMYCIN, DAPTOMYCIN
 DAUNORUBICIN HYDROCHLORIDE, DAUNORUBICIN HYDROCHLORIDE
 DEFEROXAMINE MESYLATE, DEFEROXAMINE MESYLATE
 DEXAMETHASONE SODIUM PHOSPHATE PRESERVATIVE FREE, DEXAMETHASONE SODIUM PHOSPHATE
 DEXAMETHASONE SODIUM PHOSPHATE, DEXAMETHASONE SODIUM PHOSPHATE
 DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE
 DEXTROSE 10% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 2.5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 5% AND POTASSIUM CHLORIDE 0.075%, DEXTROSE
 DEXTROSE 5% AND POTASSIUM CHLORIDE 0.15%, DEXTROSE
 DEXTROSE 5% AND SODIUM CHLORIDE 0.225%, DEXTROSE
 DEXTROSE 5% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 5% AND SODIUM CHLORIDE 0.45%, DEXTROSE
 DEXTROSE 5% AND SODIUM CHLORIDE 0.9%, DEXTROSE
 DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
 DIAZEPAM, DIAZEPAM
 DICYCLOMINE HYDROCHLORIDE, DICYCLOMINE HYDROCHLORIDE
 DILAUDID, HYDROMORPHONE HYDROCHLORIDE
 DIMENHYDRINATE, DIMENHYDRINATE
 DIPHENHYDRAMINE HYDROCHLORIDE PRESERVATIVE FREE, DIPHENHYDRAMINE HYDROCHLORIDE
 DIPRIVAN, PROPOFOL
 DOXORUBICIN HYDROCHLORIDE, DOXORUBICIN HYDROCHLORIDE
 DOXY 100, DOXYCYCLINE HYCLATE
 DOXY 200, DOXYCYCLINE HYCLATE
 EPHEDRINE SULFATE, EPHEDRINE SULFATE
 EPINEPHRINE, EPINEPHRINE
 ETOPOSIDE, ETOPOSIDE
 FAMOTIDINE PRESERVATIVE FREE, FAMOTIDINE
 FAMOTIDINE, FAMOTIDINE
 FENTANYL CITRATE PRESERVATIVE FREE, FENTANYL CITRATE
 FENTANYL CITRATE, FENTANYL CITRATE
 FLOXURIDINE, FLOXURIDINE
 FLUCONAZOLE IN SODIUM CHLORIDE 0.9%, FLUCONAZOLE
 FLUDARABINE PHOSPHATE, FLUDARABINE PHOSPHATE
 FLUMAZENIL, FLUMAZENIL
 FLUOROURACIL, FLUOROURACIL
 FLUPHENAZINE DECANOATE, FLUPHENAZINE DECANOATE
 FLUPHENAZINE HYDROCHLORIDE, FLUPHENAZINE HYDROCHLORIDE
 FOLIC ACID, FOLIC ACID
 FOSAPREPITANT DIMEGLUMINE, FOSAPREPITANT DIMEGLUMINE
 FOSCARNET SODIUM, FOSCARNET SODIUM
 FOSPHENYTOIN SODIUM, FOSPHENYTOIN SODIUM
 FULVESTRANT, FULVESTRANT
 FUROSEMIDE, FUROSEMIDE
 GANCICLOVIR SODIUM, GANCICLOVIR SODIUM
 GEMCITABINE HYDROCHLORIDE, GEMCITABINE HYDROCHLORIDE
 GENTAMICIN SULFATE, GENTAMICIN SULFATE
 GLUCAGON, GLUCAGON HYDROCHLORIDE
 GLYCOPYRROLATE, GLYCOPYRROLATE
 GRANISETRON HYDROCHLORIDE PRESERVATIVE FREE, GRANISETRON HYDROCHLORIDE
 GRANISETRON HYDROCHLORIDE, GRANISETRON HYDROCHLORIDE
 HALOPERIDOL DECANOATE, HALOPERIDOL DECANOATE
 HALOPERIDOL, HALOPERIDOL LACTATE
 HEPARIN SODIUM 1,000 UNITS IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, HEPARIN
 HEPARIN SODIUM 2,000 UNITS IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, HEPARIN
 HEPARIN SODIUM 25,000 UNITS IN DEXTROSE 5%, HEPARIN SODIUM
 HEPARIN SODIUM 25,000 UNITS IN SODIUM CHLORIDE 0.45%, HEPARIN SODIUM
 HEPARIN SODIUM IN PLASTIC CONTAINER, HEPARIN SODIUM
 HEPARIN SODIUM PRESERVATIVE FREE, HEPARIN SODIUM
 HEPARIN SODIUM, HEPARIN SODIUM
 HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
 ICATIBANT ACETATE, ICATIBANT ACETATE
 IFOSFAMIDE, IFOSFAMIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** F ****

* FRESENIUS KABI USA LLC
 INDOMETHACIN, INDOMETHACIN
 IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE
 KABIVEN IN PLASTIC CONTAINER, AMINO ACIDS
 KETAMINE HYDROCHLORIDE, KETAMINE HYDROCHLORIDE
 KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
 LACOSAMIDE, LACOSAMIDE
 LACTATED RINGER'S IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 LEUCOVORIN CALCIUM PRESERVATIVE FREE, LEUCOVORIN CALCIUM
 LEUCOVORIN CALCIUM, LEUCOVORIN CALCIUM
 LEVETIRACETAM, LEVETIRACETAM
 LEVOFLOXACIN IN DEXTROSE 5% IN PLASTIC CONTAINER, LEVOFLOXACIN
 LEVOTHYROXINE SODIUM, LEVOTHYROXINE SODIUM
 LIDOCAINE HYDROCHLORIDE IN PLASTIC CONTAINER, LIDOCAINE HYDROCHLORIDE
 LIDOCAINE HYDROCHLORIDE PRESERVATIVE FREE, LIDOCAINE HYDROCHLORIDE
 LIDOCAINE HYDROCHLORIDE, LIDOCAINE HYDROCHLORIDE
 LINEZOLID, LINEZOLID
 LORAZEPAM, LORAZEPAM
 MAGNESIUM SULFATE IN DEXTROSE 5% IN PLASTIC CONTAINER, MAGNESIUM SULFATE
 MAGNESIUM SULFATE IN PLASTIC CONTAINER, MAGNESIUM SULFATE
 MAGNESIUM SULFATE, MAGNESIUM SULFATE
 MANNITOL 25%, MANNITOL
 MELPHALAN HYDROCHLORIDE, MELPHALAN HYDROCHLORIDE
 MESNA, MESNA
 METHOCARBAMOL, METHOCARBAMOL
 METHOTREXATE PRESERVATIVE FREE, METHOTREXATE SODIUM
 METHOTREXATE SODIUM, METHOTREXATE SODIUM
 METHYLPREDNISOLONE SODIUM SUCCINATE, METHYLPREDNISOLONE SODIUM SUCCINATE
 METOCLOPRAMIDE HYDROCHLORIDE, METOCLOPRAMIDE HYDROCHLORIDE
 METOPROLOL TARTRATE, METOPROLOL TARTRATE
 MICAFUNGIN SODIUM, MICAFUNGIN SODIUM
 MIDAZOLAM HYDROCHLORIDE PRESERVATIVE FREE, MIDAZOLAM HYDROCHLORIDE
 MIDAZOLAM HYDROCHLORIDE, MIDAZOLAM HYDROCHLORIDE
 MILRINONE LACTATE, MILRINONE LACTATE
 MITOMYCIN, MITOMYCIN
 MITOXANTRONE HYDROCHLORIDE, MITOXANTRONE HYDROCHLORIDE
 MORPHINE SULFATE, MORPHINE SULFATE
 MOXIFLOXACIN HYDROCHLORIDE, MOXIFLOXACIN HYDROCHLORIDE
 MULTIPLE ELECTROLYTES INJECTION TYPE 1 USP PH 5.5, MAGNESIUM CHLORIDE
 MULTIPLE ELECTROLYTES INJECTION TYPE 1 USP PH 7.4, MAGNESIUM CHLORIDE
 NALOXONE HYDROCHLORIDE, NALOXONE HYDROCHLORIDE
 NAROPIN, ROPIVACAINE HYDROCHLORIDE
 NEBUPENT, PENTAMIDINE ISETHIONATE
 NEOSTIGMINE METHYLSULFATE, NEOSTIGMINE METHYLSULFATE
 NESACAINE, CHLOROPROCAINE HYDROCHLORIDE
 NESACAINE-MPF, CHLOROPROCAINE HYDROCHLORIDE
 NOREPINEPHRINE BITARTRATE, NOREPINEPHRINE BITARTRATE
 OCTREOTIDE ACETATE (PRESERVATIVE FREE), OCTREOTIDE ACETATE
 OCTREOTIDE ACETATE, OCTREOTIDE ACETATE
 OMEGAVEN, FISH OIL TRIGLYCERIDES
 ONDANSETRON HYDROCHLORIDE PRESERVATIVE FREE, ONDANSETRON HYDROCHLORIDE
 ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
 OXACILLIN SODIUM, OXACILLIN SODIUM
 OXALIPLATIN, OXALIPLATIN
 OXYTOCIN, OXYTOCIN
 PACLITAXEL, PACLITAXEL
 PALONOSETRON HYDROCHLORIDE, PALONOSETRON HYDROCHLORIDE
 PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
 PEMETREXED DISODIUM, PEMETREXED DISODIUM
 PENTAM, PENTAMIDINE ISETHIONATE
 PERIKABIVEN IN PLASTIC CONTAINER, AMINO ACIDS
 PHENYLEPHRINE HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE
 POLOCAINE, MEPIVACAINE HYDROCHLORIDE
 POLOCAINE-MPF, MEPIVACAINE HYDROCHLORIDE
 POLYMYXIN B SULFATE, POLYMYXIN B SULFATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** F ****

* FRESENIUS KABI USA LLC
 POSACONAZOLE, POSACONAZOLE
 POTASSIUM ACETATE, POTASSIUM ACETATE
 POTASSIUM CHLORIDE 0.15% IN SODIUM CHLORIDE 0.45%, POTASSIUM CHLORIDE
 POTASSIUM CHLORIDE 0.15% IN SODIUM CHLORIDE 0.9%, POTASSIUM CHLORIDE
 POTASSIUM CHLORIDE 0.3% AND SODIUM CHLORIDE 0.9%, POTASSIUM CHLORIDE
 POTASSIUM CHLORIDE 10MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.225% IN PLASTIC
 POTASSIUM CHLORIDE 10MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45%, DEXTROSE
 POTASSIUM CHLORIDE 10MEQ, POTASSIUM CHLORIDE
 POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.225% IN PLASTIC
 POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45%, DEXTROSE
 POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9%, DEXTROSE
 POTASSIUM CHLORIDE 20MEQ, POTASSIUM CHLORIDE
 POTASSIUM CHLORIDE 30MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45%, DEXTROSE
 POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45%, DEXTROSE
 POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9%, DEXTROSE
 POTASSIUM CHLORIDE 40MEQ, POTASSIUM CHLORIDE
 POTASSIUM CHLORIDE IN PLASTIC CONTAINER, POTASSIUM CHLORIDE
 POTASSIUM PHOSPHATES, POTASSIUM PHOSPHATE, DIBASIC
 PROGESTERONE, PROGESTERONE
 PROPRANOLOL HYDROCHLORIDE, PROPRANOLOL HYDROCHLORIDE
 PROTAMINE SULFATE, PROTAMINE SULFATE
 PYRIDOXINE HYDROCHLORIDE, PYRIDOXINE HYDROCHLORIDE
 REMIFENTANIL HYDROCHLORIDE, REMIFENTANIL HYDROCHLORIDE
 RIFAMPIN, RIFAMPIN
 ROCURONIUM BROMIDE, ROCURONIUM BROMIDE
 ROMIDEPSIN, ROMIDEPSIN
 SENSORCAINE, BUPIVACAINE HYDROCHLORIDE
 SENSORCAINE, BUPIVACAINE HYDROCHLORIDE
 SMOFLIPID 20%, FISH OIL
 SODIUM ACETATE, SODIUM ACETATE
 SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, SODIUM CHLORIDE
 SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, SODIUM CHLORIDE
 SODIUM CHLORIDE 14.6%, SODIUM CHLORIDE
 SODIUM CHLORIDE 23.4%, SODIUM CHLORIDE
 SODIUM CHLORIDE 3% IN PLASTIC CONTAINER, SODIUM CHLORIDE
 SODIUM PHOSPHATES, SODIUM PHOSPHATE, DIBASIC, HEPTAHYDRATE
 STERILE WATER FOR INJECTION IN PLASTIC CONTAINER, STERILE WATER FOR INJECTION
 STERILE WATER FOR INJECTION, STERILE WATER FOR INJECTION
 STERILE WATER FOR IRRIGATION, STERILE WATER FOR IRRIGATION
 SUCCINYLCHOLINE CHLORIDE, SUCCINYLCHOLINE CHLORIDE
 SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
 TERBUTALINE SULFATE, TERBUTALINE SULFATE
 THIAMINE HYDROCHLORIDE, THIAMINE HYDROCHLORIDE
 TIGECYCLINE, TIGECYCLINE
 TOBRAMYCIN SULFATE (PHARMACY BULK), TOBRAMYCIN SULFATE
 TOBRAMYCIN SULFATE, TOBRAMYCIN SULFATE
 TOPOTECAN HYDROCHLORIDE, TOPOTECAN HYDROCHLORIDE
 TRANEXAMIC ACID, TRANEXAMIC ACID
 VALPROATE SODIUM, VALPROATE SODIUM
 VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE
 VASOPRESSIN, VASOPRESSIN
 VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE
 VIBISONE, CYANOCOBALAMIN
 VINBLASTINE SULFATE, VINBLASTINE SULFATE
 XYLOCAINE W/ EPINEPHRINE, EPINEPHRINE
 XYLOCAINE, LIDOCAINE HYDROCHLORIDE
 ZINC SULFATE, ZINC SULFATE
 ZOLEDRONIC ACID, ZOLEDRONIC ACID

FRESENIUS MEDCL

* FRESENIUS MEDICAL CARE NORTH AMERICA
 DELFLEX W/ DEXTROSE 1.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 DELFLEX W/ DEXTROSE 1.5% LOW MAGNESIUM IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 DELFLEX W/ DEXTROSE 1.5% LOW MAGNESIUM LOW CALCIUM IN PLASTIC CONTAINER, CALCIUM
 DELFLEX W/ DEXTROSE 2.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** F ****

- * FRESENIUS MEDICAL CARE NORTH AMERICA
 - DELFLEX W/ DEXTROSE 2.5% LOW MAGNESIUM IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 - DELFLEX W/ DEXTROSE 2.5% LOW MAGNESIUM LOW CALCIUM IN PLASTIC CONTAINER, CALCIUM
 - DELFLEX W/ DEXTROSE 4.25% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 - DELFLEX W/ DEXTROSE 4.25% LOW MAGNESIUM IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 - DELFLEX W/ DEXTROSE 4.25% LOW MAGNESIUM LOW CALCIUM IN PLASTIC CONTAINER, CALCIUM
 - SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, SODIUM CHLORIDE

**** G ******GALDERMA LABS**

- * GALDERMA LABORATORIES INC
 - CLOBEX, CLOBETASOL PROPIONATE
 - EPIDUO FORTE, ADAPALENE

GALDERMA LABS LP

- * GALDERMA LABORATORIES L P
 - CLOBEX, CLOBETASOL PROPIONATE
- * GALDERMA LABORATORIES LP
 - AKLIEF, TRIFAROTENE
 - CLOBEX, CLOBETASOL PROPIONATE
 - DESOWEN, DESONIDE
 - DIFFERIN, ADAPALENE
 - DIFFERIN, ADAPALENE (OTC)
 - EPIDUO, ADAPALENE
 - EPSOLAY, BENZOYL PEROXIDE
 - METROCREAM, METRONIDAZOLE
 - METROGEL, METRONIDAZOLE
 - METROLOTION, METRONIDAZOLE
 - MIRVASO, BRIMONIDINE TARTRATE
 - ORACEA, DOXYCYCLINE
 - SOOLANTRA, IVERMECTIN
 - TRI-LUMA, FLUOCINOLONE ACETONIDE
 - TWYNEO, BENZOYL PEROXIDE
 - VECTICAL, CALCITRIOL

GALT PHARMS

- * GALT PHARMACEUTICALS LLC
 - DORAL, QUAZEPAM
 - ORAVIG, MICONAZOLE
 - ORPHENGESIC FORTE, ASPIRIN
 - ORPHENGESIC, ASPIRIN

GE HEALTHCARE

- * GE HEALTHCARE
 - ADREVIEW, IOBENGUANE SULFATE I-123
 - CERETEC, TECHNETIUM TC-99M EXAMETAZIME KIT
 - CERIANNA, FLUOROESTRADIOL F-18
 - CLARISCAN, GADOTERATE MEGLUMINE
 - INDIUM IN 111 OXYQUINOLINE, INDIUM IN-111 OXYQUINOLINE
 - MPI INDIUM DTPA IN 111, INDIUM IN-111 PENTETATE DISODIUM
 - OMNIPAQUE 12, IOHEXOL
 - OMNIPAQUE 140, IOHEXOL
 - OMNIPAQUE 180, IOHEXOL
 - OMNIPAQUE 240, IOHEXOL
 - OMNIPAQUE 300, IOHEXOL
 - OMNIPAQUE 350, IOHEXOL
 - OMNIPAQUE 9, IOHEXOL
 - OMNISCAN, GADODIAMIDE
 - OPTISON, ALBUMIN HUMAN
 - REGADENOSON, REGADENOSON
 - VISIPAQUE 270, IODIXANOL
 - VISIPAQUE 320, IODIXANOL
 - VIZAMYL, FLUTEMETAMOL F-18

GE HLTHCARE

- * GE HEALTHCARE INC
 - FLYRCADO, FLURPIRIDAZ F-18

GE HLTHCARE INC

- * GE HEALTHCARE INC

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** G ****

* GE HEALTHCARE INC

DATSCAN, IOFLUPANE I-123

GENBIOPRO

* GENBIOPRO INC

MIFEPRISTONE, MIFEPRISTONE

GENENTECH

* GENENTECH INC

ERIVEDGE, VISMODEGIB

GENENTECH INC

* GENENTECH INC

COTELLIC, COBIMETINIB FUMARATE

ESBRIET, PIRFENIDONE

EVRYSDI, RISDIPLAM

ITOVEBI, INAVOLISIB

ROZLYTREK, ENTRECTINIB

XOFLUZA, BALOXAVIR MARBOXIL

GENUS

* GENUS LIFESCIENCES INC

AMCINONIDE, AMCINONIDE

CARBINOXAMINE MALEATE, CARBINOXAMINE MALEATE

CHLORPROMAZINE HYDROCHLORIDE, CHLORPROMAZINE HYDROCHLORIDE

CLEMASTINE FUMARATE, CLEMASTINE FUMARATE

FLURBIPROFEN, FLURBIPROFEN

HYCODAN, HOMATROPINE METHYLBROMIDE

HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN

ISONIAZID, ISONIAZID

ISOSORBIDE MONONITRATE, ISOSORBIDE MONONITRATE

MEPERIDINE HYDROCHLORIDE, MEPERIDINE HYDROCHLORIDE

METOCLOPRAMIDE HYDROCHLORIDE, METOCLOPRAMIDE HYDROCHLORIDE

NYSTATIN, NYSTATIN

PHENYLEPHRINE HYDROCHLORIDE AND PROMETHAZINE HYDROCHLORIDE, PHENYLEPHRINE

POKONZA, POTASSIUM CHLORIDE

POTASSIUM CHLORIDE, POTASSIUM CHLORIDE

PROMETH HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE W/CODEINE PHOSPHATE, CODEINE

GENUS LIFESCIENCES

* GENUS LIFE SCIENCES INC

ACETAMINOPHEN AND CODEINE PHOSPHATE, ACETAMINOPHEN

AMCINONIDE, AMCINONIDE

LEVOLET, LEVOTHYROXINE SODIUM **

METAPROTERENOL SULFATE, METAPROTERENOL SULFATE

OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE

POTASSIUM CHLORIDE, POTASSIUM CHLORIDE

GENZYME

* GENZYME CORP

CLOLAR, CLOFARABINE

MOZOBIL, PLERIXAFOR

RENAGEL, SEVELAMER HYDROCHLORIDE

REVELA, SEVELAMER CARBONATE

GENZYME CORP

* GENZYME CORP

CAPRELSA, VANDETANIB

CERDELGA, ELIGLUSTAT TARTRATE

GERON

* GERON CORP

RYTELO, IMETELSTAT SODIUM

GILEAD

* GILEAD SCIENCES INC

CAYSTON, AZTREONAM

EMTRIVA, EMTRICITABINE

LETAIRIS, AMBRISENTAN

TRUVADA, EMTRICITABINE

GILEAD SCIENCES INC

* GILEAD SCIENCES INC

BIKTARVY, BICTEGRAVIR SODIUM

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** G ****

* GILEAD SCIENCES INC
 COMPLERA, EMTRICITABINE
 DESCOVY, EMTRICITABINE
 EPCLUSA, SOFOSBUVIR
 GENVOYA, COBICISTAT
 HARVONI, LEDIPASVIR
 LIVDELZI, SELADELPAVIR LYSINE
 ODEFSEY, EMTRICITABINE
 SOVALDI, SOFOSBUVIR
 STRIBILD, COBICISTAT
 SUNLENCA, LENACAPAVIR SODIUM
 TYBOST, COBICISTAT
 VEKLURY, REMDESIVIR
 VEMLIDY, TENOFOVIR ALAFENAMIDE FUMARATE
 VIREAD, TENOFOVIR DISOPROXIL FUMARATE
 VOSEVI, SOFOSBUVIR
 ZYDELIG, IDELALISIB

GISKIT

* GISKIT PHARMA BV
 EXEM FOAM KIT, AIR POLYMER-TYPE A

GLAND

* GLAND PHARMA LTD
 CASPOFUNGIN ACETATE, CASPOFUNGIN ACETATE
 CETRORELIX ACETATE, CETRORELIX ACETATE
 DECITABINE, DECITABINE
 DESMOPRESSIN ACETATE, DESMOPRESSIN ACETATE
 DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE
 DEXRAZOXANE HYDROCHLORIDE, DEXRAZOXANE HYDROCHLORIDE
 ENOXAPARIN SODIUM (PRESERVATIVE FREE), ENOXAPARIN SODIUM
 HEPARIN SODIUM, HEPARIN SODIUM
 KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
 LATANOPROST, LATANOPROST
 MEROPENEM, MEROPENEM
 MESNA, MESNA
 OXALIPLATIN, OXALIPLATIN
 POLYMYXIN B SULFATE, POLYMYXIN B SULFATE
 TRANEXAMIC ACID, TRANEXAMIC ACID
 VECURONIUM BROMIDE, VECURONIUM BROMIDE
 ZOLEDRONIC ACID, ZOLEDRONIC ACID

GLAND PHARMA LTD

* GLAND PHARMA LTD
 ADENOSINE, ADENOSINE
 ALCAFTADINE, ALCAFTADINE (OTC)
 ALLOPURINOL SODIUM, ALLOPURINOL SODIUM
 AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE
 ARGATROBAN IN SODIUM CHLORIDE, ARGATROBAN
 ARGATROBAN, ARGATROBAN
 ARSENIC TRIOXIDE, ARSENIC TRIOXIDE
 AZELASTINE HYDROCHLORIDE, AZELASTINE HYDROCHLORIDE
 AZITHROMYCIN, AZITHROMYCIN
 BIMATOPROST, BIMATOPROST
 BROMFENAC SODIUM, BROMFENAC SODIUM
 BUMETANIDE, BUMETANIDE
 CALCITRIOL, CALCITRIOL
 CARBOPLATIN, CARBOPLATIN
 CARBOPROST TROMETHAMINE, CARBOPROST TROMETHAMINE
 CHLOROTHIAZIDE SODIUM, CHLOROTHIAZIDE SODIUM
 CHLORPROMAZINE HYDROCHLORIDE, CHLORPROMAZINE HYDROCHLORIDE
 CISPLATIN, CISPLATIN
 CLOFARABINE, CLOFARABINE
 CYANOCOBALAMIN, CYANOCOBALAMIN
 CYTARABINE, CYTARABINE
 DEFEROXAMINE MESYLATE, DEFEROXAMINE MESYLATE
 DESMOPRESSIN ACETATE, DESMOPRESSIN ACETATE
 DEXAMETHASONE SODIUM PHOSPHATE, DEXAMETHASONE SODIUM PHOSPHATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** G ******* GLAND PHARMA LTD**

DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE
 DIHYDROERGOTAMINE MESYLATE, DIHYDROERGOTAMINE MESYLATE
 DIPHENHYDRAMINE HYDROCHLORIDE, DIPHENHYDRAMINE HYDROCHLORIDE
 DOCETAXEL, DOCETAXEL
 DORZOLAMIDE HYDROCHLORIDE AND TIMOLOL MALEATE, DORZOLAMIDE HYDROCHLORIDE
 DORZOLAMIDE HYDROCHLORIDE, DORZOLAMIDE HYDROCHLORIDE
 DOXERCALCIFEROL, DOXERCALCIFEROL
 DOXORUBICIN HYDROCHLORIDE, DOXORUBICIN HYDROCHLORIDE
 DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
 EDARAVONE, EDARAVONE
 EPHEDRINE SULFATE, EPHEDRINE SULFATE
 ERIBULIN MESYLATE, ERIBULIN MESYLATE
 ERTAPENEM SODIUM, ERTAPENEM SODIUM
 ERYTHROMYCIN LACTOBIONATE, ERYTHROMYCIN LACTOBIONATE
 ESMOLOL HYDROCHLORIDE, ESMOLOL HYDROCHLORIDE
 ESOMEPRAZOLE SODIUM, ESOMEPRAZOLE SODIUM
 ETOMIDATE, ETOMIDATE
 FLUOROURACIL, FLUOROURACIL
 FLUPHENAZINE DECANOATE, FLUPHENAZINE DECANOATE
 FOMEPIZOLE, FOMEPIZOLE
 FOSCARNET SODIUM, FOSCARNET SODIUM
 FOSPHENYTOIN SODIUM, FOSPHENYTOIN SODIUM
 FUROSEMIDE, FUROSEMIDE
 GANIRELIX ACETATE, GANIRELIX ACETATE
 GEMCITABINE HYDROCHLORIDE, GEMCITABINE HYDROCHLORIDE
 GLYCOPYRROLATE, GLYCOPYRROLATE
 HALOPERIDOL DECANOATE, HALOPERIDOL DECANOATE
 HALOPERIDOL, HALOPERIDOL LACTATE
 IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE
 ISOPROTERENOL HYDROCHLORIDE, ISOPROTERENOL HYDROCHLORIDE
 KETAMINE HYDROCHLORIDE, KETAMINE HYDROCHLORIDE
 LABETALOL HYDROCHLORIDE, LABETALOL HYDROCHLORIDE
 LACOSAMIDE, LACOSAMIDE
 LEVETIRACETAM IN SODIUM CHLORIDE, LEVETIRACETAM
 LEVOFLOXACIN IN DEXTROSE 5% IN PLASTIC CONTAINER, LEVOFLOXACIN
 LEVOFLOXACIN, LEVOFLOXACIN
 LEVOLEUCOVORIN CALCIUM, LEVOLEUCOVORIN CALCIUM
 LINCOMYCIN HYDROCHLORIDE, LINCOMYCIN HYDROCHLORIDE
 MAGNESIUM SULFATE IN PLASTIC CONTAINER, MAGNESIUM SULFATE
 MELPHALAN HYDROCHLORIDE, MELPHALAN HYDROCHLORIDE
 METHOCARBAMOL, METHOCARBAMOL
 METOPROLOL TARTRATE, METOPROLOL TARTRATE
 METRONIDAZOLE IN PLASTIC CONTAINER, METRONIDAZOLE
 MIDAZOLAM HYDROCHLORIDE, MIDAZOLAM HYDROCHLORIDE
 MIDAZOLAM IN 0.9% SODIUM CHLORIDE, MIDAZOLAM
 MILRINONE LACTATE IN DEXTROSE 5% IN PLASTIC CONTAINER, MILRINONE LACTATE
 MITOMYCIN, MITOMYCIN
 MOXIFLOXACIN HYDROCHLORIDE, MOXIFLOXACIN HYDROCHLORIDE
 NELARABINE, NELARABINE
 NEOSTIGMINE METHYLSULFATE, NEOSTIGMINE METHYLSULFATE
 NOREPINEPHRINE BITARTRATE, NOREPINEPHRINE BITARTRATE
 OCTREOTIDE ACETATE, OCTREOTIDE ACETATE
 OLOPATADINE HYDROCHLORIDE, OLOPATADINE HYDROCHLORIDE (OTC)
 ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
 OXALIPLATIN, OXALIPLATIN
 PACLITAXEL, PACLITAXEL
 PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
 PHENYLEPHRINE HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE
 PHYTONADIONE, PHYTONADIONE
 PLERIXAFOR, PLERIXAFOR
 POSACONAZOLE, POSACONAZOLE
 PROCAINAMIDE HYDROCHLORIDE, PROCAINAMIDE HYDROCHLORIDE
 PROCHLORPERAZINE EDISYLATE, PROCHLORPERAZINE EDISYLATE
 REGADENOSON, REGADENOSON

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** G ****

* GLAND PHARMA LTD
 ROCURONIUM BROMIDE, ROCURONIUM BROMIDE
 ROPIVACAINE HYDROCHLORIDE, ROPIVACAINE HYDROCHLORIDE
 SUCCINYLCHOLINE CHLORIDE, SUCCINYLCHOLINE CHLORIDE
 TEMSIROLIMUS, TEMSIROLIMUS
 THIOTEPA, THIOTEPA
 TIMOLOL MALEATE, TIMOLOL MALEATE
 TIROFIBAN HYDROCHLORIDE, TIROFIBAN HYDROCHLORIDE
 TOBRAMYCIN SULFATE (PHARMACY BULK), TOBRAMYCIN SULFATE
 TOBRAMYCIN SULFATE, TOBRAMYCIN SULFATE
 TOBRAMYCIN, TOBRAMYCIN
 TRANEXAMIC ACID, TRANEXAMIC ACID
 TRAVOPROST, TRAVOPROST
 VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE
 VASOPRESSIN, VASOPRESSIN
 VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE
 VORICONAZOLE, VORICONAZOLE
 ZINC SULFATE, ZINC SULFATE
 ZIPRASIDONE MESYLATE, ZIPRASIDONE MESYLATE
 ZOLEDRONIC ACID, ZOLEDRONIC ACID
 ZOLEDRONIC, ZOLEDRONIC ACID

GLAUKOS

* GLAUKOS CORP
 IDOSE TR, TRAVOPROST
 PHOTREXA VISCOUS IN DEXTRAN 20%, RIBOFLAVIN 5'-PHOSPHATE
 PHOTREXA, RIBOFLAVIN 5'-PHOSPHATE

GLAXO GRP ENGLAND

* GLAXO GROUP LTD ENGLAND DBA GLAXOSMITHKLINE
 INCRUSE ELLIPTA, UMECLIDINIUM BROMIDE

GLAXO GRP LTD

* GLAXO GROUP LTD DBA GLAXOSMITHKLINE
 FLOVENT HFA, FLUTICASONE PROPIONATE
 * GLAXO GROUP LTD ENGLAND DBA GLAXOSMITHKLINE
 ADVAIR DISKUS 100/50, FLUTICASONE PROPIONATE
 ADVAIR DISKUS 250/50, FLUTICASONE PROPIONATE
 ADVAIR DISKUS 500/50, FLUTICASONE PROPIONATE
 ADVAIR HFA, FLUTICASONE PROPIONATE
 BREO ELLIPTA, FLUTICASONE FUROATE
 FLOVENT DISKUS 100, FLUTICASONE PROPIONATE
 FLOVENT DISKUS 250, FLUTICASONE PROPIONATE
 FLOVENT DISKUS 50, FLUTICASONE PROPIONATE

GLAXOSMITHKLINE

* GLAXOSMITHKLINE
 IMITREX STATDOSE, SUMATRIPTAN SUCCINATE
 IMITREX, SUMATRIPTAN
 IMITREX, SUMATRIPTAN SUCCINATE
 MALARONE PEDIATRIC, ATOVAQUONE
 MALARONE, ATOVAQUONE
 RELENZA, ZANAMIVIR
 VALTREX, VALACYCLOVIR HYDROCHLORIDE
 WELLBUTRIN SR, BUPROPION HYDROCHLORIDE
 * GLAXOSMITHKLINE INTELLECTUAL PROPERTY DEVELOPMENT LTD ENGLAND
 ANORO ELLIPTA, UMECLIDINIUM BROMIDE
 ARNUITY ELLIPTA, FLUTICASONE FUROATE
 KRINTAFEL, TAFENOQUINE SUCCINATE
 TRELEGY ELLIPTA, FLUTICASONE FUROATE
 * GLAXOSMITHKLINE INTELLECTUAL PROPERTY LTD ENGLAND
 SEREVENT, SALMETEROL XINAFOATE
 VENTOLIN HFA, ALBUTEROL SULFATE
 * GLAXOSMITHKLINE LLC
 OJJAARA, MOMELOTINIB DIHYDROCHLORIDE
 ZEJULA, NIRAPARIB TOSYLATE

GLAXOSMITHKLINE LLC

* GLAXOSMITHKLINE LLC
 FLOLAN, EPOPROSTENOL SODIUM

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** G ******* GLAXOSMITHKLINE LLC**

LAMICTAL CD, LAMOTRIGINE
 LAMICTAL ODT, LAMOTRIGINE
 LAMICTAL XR, LAMOTRIGINE
 LAMICTAL, LAMOTRIGINE
 MEPRON, ATOVAQUONE

GLENMARK PHARMS*** GLENMARK PHARMACEUTICALS INC USA**

CICLOPIROX, CICLOPIROX
 CLOTRIMAZOLE, CLOTRIMAZOLE
 MUPIROCIN, MUPIROCIN

*** GLENMARK PHARMACEUTICALS LTD**

MOEXIPRIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE

GLENMARK PHARMS INC*** GLENMARK PHARMACEUTICALS INC USA**

ACETYLCYSTEINE, ACETYLCYSTEINE
 ADAPALENE, ADAPALENE (OTC)
 BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
 CALCIPOTRIENE, CALCIPOTRIENE
 CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 CICLOPIROX, CICLOPIROX
 FAMOTIDINE, FAMOTIDINE (OTC)
 FULVESTRANT, FULVESTRANT
 IMIQUIMOD, IMIQUIMOD
 LANSOPRAZOLE, LANSOPRAZOLE (OTC)
 LITHIUM CARBONATE, LITHIUM CARBONATE
 MOMETASONE FUROATE, MOMETASONE FUROATE
 MUPIROCIN, MUPIROCIN CALCIUM
 NIZATIDINE, NIZATIDINE
 OLOPATADINE HYDROCHLORIDE, OLOPATADINE HYDROCHLORIDE (OTC)
 RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
 ZONISAMIDE, ZONISAMIDE

GLENMARK PHARMS LTD*** GLENMARK PHARMACEUTICALS LTD**

ACETAMINOPHEN AND IBUPROFEN, ACETAMINOPHEN (OTC)
 ADAPALENE AND BENZOYL PEROXIDE, ADAPALENE
 ALCLOMETASONE DIPROPIONATE, ALCLOMETASONE DIPROPIONATE
 ALYACEN 1/35, ETHINYL ESTRADIOL
 ALYACEN 7/7/7, ETHINYL ESTRADIOL
 APREMILAST, APREMILAST
 ASHLYNA, ETHINYL ESTRADIOL
 ATOMOXETINE HYDROCHLORIDE, ATOMOXETINE HYDROCHLORIDE
 ATOVAQUONE AND PROGUANIL HYDROCHLORIDE, ATOVAQUONE
 BISOPROLOL FUMARATE AND HYDROCHLOROTHIAZIDE, BISOPROLOL FUMARATE
 BRIELLYN, ETHINYL ESTRADIOL
 BRIMONIDINE TARTRATE AND TIMOLOL MALEATE, BRIMONIDINE TARTRATE
 CARVEDILOL, CARVEDILOL
 CHLORPROMAZINE HYDROCHLORIDE, CHLORPROMAZINE HYDROCHLORIDE
 CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE
 CLINDAMYCIN HYDROCHLORIDE, CLINDAMYCIN HYDROCHLORIDE
 CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
 CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
 COLESEVELAM HYDROCHLORIDE, COLESEVELAM HYDROCHLORIDE
 DESMOPRESSIN ACETATE, DESMOPRESSIN ACETATE
 DESONIDE, DESONIDE
 DESOXIMETASONE, DESOXIMETASONE
 DICLOFENAC SODIUM, DICLOFENAC SODIUM
 DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
 DIMETHYL FUMARATE, DIMETHYL FUMARATE
 DROSPIRENONE AND ETHINYL ESTRADIOL, DROSPIRENONE
 ESTRADIOL, ESTRADIOL
 ESZOPICLONE, ESZOPICLONE
 EZETIMIBE AND SIMVASTATIN, EZETIMIBE
 EZETIMIBE, EZETIMIBE
 FELODIPINE, FELODIPINE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** G **

* GLENMARK PHARMACEUTICALS LTD
FENOFIBRATE (MICRONIZED), FENOFIBRATE
FINGOLIMOD HYDROCHLORIDE, FINGOLIMOD HYDROCHLORIDE
FLUCINOLONE ACETONIDE, FLUOCINOLONE ACETONIDE
FLUCONAZOLE, FLUCONAZOLE
FLUOCINONIDE ACETONIDE, FLUOCINOLONE ACETONIDE
FLUOCINONIDE, FLUOCINONIDE
FLUPHENAZINE HYDROCHLORIDE, FLUPHENAZINE HYDROCHLORIDE
FLUTICASONE PROPIONATE, FLUTICASONE PROPIONATE
FROVATRIPTAN SUCCINATE, FROVATRIPTAN SUCCINATE
GABAPENTIN, GABAPENTIN
HAILEY 24 FE, ETHINYL ESTRADIOL
HAILEY FE 1/20, ETHINYL ESTRADIOL
HEATHER, NORETHINDRONE
HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
HYDROCORTISONE BUTYRATE, HYDROCORTISONE BUTYRATE
HYDROCORTISONE VALERATE, HYDROCORTISONE VALERATE
INDOMETHACIN, INDOMETHACIN
LACOSAMIDE, LACOSAMIDE
LAMOTRIGINE, LAMOTRIGINE
LEVOCETIRIZINE DIHYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE
LEVOFLOXACIN, LEVOFLOXACIN
LEVONORGESTREL AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
LEVONORGESTREL, LEVONORGESTREL (OTC)
LIDOCAINE, LIDOCAINE
LITHIUM CARBONATE, LITHIUM CARBONATE
MARLISSA, ETHINYL ESTRADIOL
MELOXICAM, MELOXICAM
METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
METRONIDAZOLE, METRONIDAZOLE
MOEXIPRIL HYDROCHLORIDE, MOEXIPRIL HYDROCHLORIDE
MOMETASONE FUROATE, MOMETASONE FUROATE
MONTELUKAST SODIUM, MONTELUKAST SODIUM
NAPROXEN SODIUM, NAPROXEN SODIUM
NAPROXEN, NAPROXEN
NEBIVOLOL HYDROCHLORIDE, NEBIVOLOL HYDROCHLORIDE
NICARDIPINE HYDROCHLORIDE, NICARDIPINE HYDROCHLORIDE
NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL AND FERROUS FUMARATE, ETHINYL ESTRADIOL
NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
NORETHINDRONE ACETATE, NORETHINDRONE ACETATE
NORETHINDRONE, NORETHINDRONE
NORGESTIMATE AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
NYSTATIN AND TRIAMCINOLONE ACETONIDE, NYSTATIN
OLMESARTAN MEDOXOMIL, OLMESARTAN MEDOXOMIL
OMEPRAZOLE, OMEPRAZOLE
ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
ONDANSETRON, ONDANSETRON
OXCARBAZEPINE, OXCARBAZEPINE
PIMECROLIMUS, PIMECROLIMUS
POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE
PRAVASTATIN SODIUM, PRAVASTATIN SODIUM
PROCHLORPERAZINE MALEATE, PROCHLORPERAZINE MALEATE
PROPAFENONE HYDROCHLORIDE, PROPAFENONE HYDROCHLORIDE
RALOXIFENE HYDROCHLORIDE, RALOXIFENE HYDROCHLORIDE
RANOLAZINE, RANOLAZINE
RILUZOLE, RILUZOLE
ROPINIROLE HYDROCHLORIDE, ROPINIROLE HYDROCHLORIDE
RUFINAMIDE, RUFINAMIDE
SAXAGLIPTIN, SAXAGLIPTIN HYDROCHLORIDE
SEVELAMER HYDROCHLORIDE, SEVELAMER HYDROCHLORIDE
SIROLIMUS, SIROLIMUS
SODIUM PHENYLBUTYRATE, SODIUM PHENYLBUTYRATE
SULFAMETHOXAZOLE AND TRIMETHOPRIM, SULFAMETHOXAZOLE
TACROLIMUS, TACROLIMUS

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** G ****

* GLENMARK PHARMACEUTICALS LTD
 TELMISARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 TELMISARTAN, TELMISARTAN
 TERBINAFFINE HYDROCHLORIDE, TERBINAFFINE HYDROCHLORIDE
 THEOPHYLLINE, THEOPHYLLINE
 TOPIRAMATE, TOPIRAMATE
 TRANDOLAPRIL AND VERAPAMIL HYDROCHLORIDE, TRANDOLAPRIL
 TRETINOIN, TRETINOIN
 TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE
 TROSPIMUM CHLORIDE, TROSPIMUM CHLORIDE
 URSODIOL, URSODIOL
 VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE
 VIORELE, DESOGESTREL
 VORICONAZOLE, VORICONAZOLE
 ZOLMITRIPTAN, ZOLMITRIPTAN

GLENMARK SPECLT

* GLENMARK SPECIALTY SA
 ABIRATERONE ACETATE, ABIRATERONE ACETATE
 ACYCLOVIR, ACYCLOVIR
 APREPITANT, APREPITANT
 ASPIRIN AND DIPYRIDAMOLE, ASPIRIN
 ATOVAQUONE, ATOVAQUONE
 AZELAIC ACID, AZELAIC ACID
 CALCIPOTRIENE, CALCIPOTRIENE
 CLINDAMYCIN PHOSPHATE AND BENZOYL PEROXIDE, BENZOYL PEROXIDE
 CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
 CLOTRIMAZOLE AND BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
 DEFERASIROX, DEFERASIROX
 DESONIDE, DESONIDE
 ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM (OTC)
 HAILEY 1.5/30, ETHINYL ESTRADIOL
 HAILEY FE 1.5/30, ETHINYL ESTRADIOL
 LINEZOLID, LINEZOLID
 NITROGLYCERIN, NITROGLYCERIN
 NORGESTIMATE AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
 ROSUVASTATIN CALCIUM, ROSUVASTATIN CALCIUM
 RYALTRIS, MOMETASONE FUROATE
 SOLIFENACIN SUCCINATE, SOLIFENACIN SUCCINATE
 TERIFLUNOMIDE, TERIFLUNOMIDE
 TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE

GLW

* GLW PHARMA GMBH
 OMEGA-3-ACID ETHYL ESTERS, OMEGA-3-ACID ETHYL ESTERS

GRANULES

* GRANULES INDIA LTD
 ACETAMINOPHEN, ACETAMINOPHEN (OTC)
 ACETAMINOPHEN, ASPIRIN AND CAFFEINE, ACETAMINOPHEN (OTC)
 CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM
 FEXOFENADINE HYDROCHLORIDE ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
 GABAPENTIN, GABAPENTIN
 GUAIFENESIN, GUAIFENESIN (OTC)
 IBUPROFEN, IBUPROFEN
 IBUPROFEN, IBUPROFEN (OTC)
 LORATADINE, LORATADINE (OTC)
 LOSARTAN POTASSIUM, LOSARTAN POTASSIUM
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 METHOCARBAMOL, METHOCARBAMOL
 METOPROLOL SUCCINATE, METOPROLOL SUCCINATE
 NAPROXEN SODIUM AND DIPHENHYDRAMINE HYDROCHLORIDE, DIPHENHYDRAMINE HYDROCHLORIDE
 NAPROXEN, NAPROXEN
 PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
 POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
 SERTRALINE HYDROCHLORIDE, SERTRALINE HYDROCHLORIDE
 TRAZODONE HYDROCHLORIDE, TRAZODONE HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** G ****

- * GRANULES INDIA LTD
 - VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE
 - ZONISAMIDE, ZONISAMIDE
- * GRANULES PHARMACEUTICALS INC
 - ACETAMINOPHEN AND IBUPROFEN, ACETAMINOPHEN (OTC)
 - AMPHETAMINE SULFATE, AMPHETAMINE SULFATE
 - BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
 - BUTALBITAL AND ACETAMINOPHEN, ACETAMINOPHEN
 - BUTALBITAL, ACETAMINOPHEN AND CAFFEINE, ACETAMINOPHEN
 - COLCHICINE, COLCHICINE
 - DEXMETHYLPHENIDATE HYDROCHLORIDE, DEXMETHYLPHENIDATE HYDROCHLORIDE
 - DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE
 - DOFETILIDE, DOFETILIDE
 - GLYCOPYRROLATE, GLYCOPYRROLATE
 - HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
 - LISDEXAMFETAMINE DIMESYLATE, LISDEXAMFETAMINE DIMESYLATE
 - METHYLERGONOVINE MALEATE, METHYLERGONOVINE MALEATE
 - METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
 - OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN
 - PENICILLAMINE, PENICILLAMINE
 - POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
 - PRazosin HYDROCHLORIDE, PRAZOSIN HYDROCHLORIDE
 - RAMELTEON, RAMELTEON
 - TROSPiUM CHLORIDE, TROSPiUM CHLORIDE
 - VALGANCICLOVIR HYDROCHLORIDE, VALGANCICLOVIR HYDROCHLORIDE

GRANULES INDIA

- * GRANULES INDIA LTD
 - IBUPROFEN, IBUPROFEN (OTC)
 - METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 - NAPROXEN SODIUM, NAPROXEN SODIUM (OTC)

GRAVITI PHARMS

- * GRAVITI PHARMACEUTICALS INC
 - VALACYCLOVIR HYDROCHLORIDE, VALACYCLOVIR HYDROCHLORIDE
- * GRAVITI PHARMACEUTICALS PRIVATE LTD
 - ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM
 - BACLOFEN, BACLOFEN
 - BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
 - CYCLOBENZAPRINE HYDROCHLORIDE, CYCLOBENZAPRINE HYDROCHLORIDE
 - ESCITALOPRAM OXALATE, ESCITALOPRAM OXALATE
 - ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM
 - ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM (OTC)
 - FAMOTIDINE, FAMOTIDINE (OTC)
 - FUROSEMIDE, FUROSEMIDE
 - HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE
 - QUETIAPINE FUMARATE, QUETIAPINE FUMARATE
 - TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE
 - TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE
 - TRAZODONE HYDROCHLORIDE, TRAZODONE HYDROCHLORIDE

GUANGZHOU NOVAKEN

- * GUANGZHOU NOVAKEN PHARMACEUTICAL CO LTD
 - ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM
 - POTASSIUM CHLORIDE, POTASSIUM CHLORIDE

GUARDIAN DRUG

- * GUARDIAN DRUG CO
 - GUAIFENESIN, GUAIFENESIN (OTC)
 - IBUPROFEN, IBUPROFEN (OTC)
 - LOPERAMIDE HYDROCHLORIDE AND SIMETHICONE, LOPERAMIDE HYDROCHLORIDE (OTC)

GUERBET

- * GUERBET
 - ELUCIREM, GADOPICLENOL
- * GUERBET LLC
 - DOTAREM, GADOTERATE MEGLUMINE
 - LIPIODOL, ETHIODIZED OIL

**** H ****

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** H ******HAEMONETICS**

* HAEMONETICS CORP
SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, SODIUM CHLORIDE

HAINAN POLY

* HAINAN POLY PHARM CO LTD
AZITHROMYCIN, AZITHROMYCIN
CISATRACURIUM BESYLATE PRESERVATIVE FREE, CISATRACURIUM BESYLATE
CISATRACURIUM BESYLATE, CISATRACURIUM BESYLATE
CYCLOPHOSPHAMIDE, CYCLOPHOSPHAMIDE
GADOTERATE MEGLUMINE, GADOTERATE MEGLUMINE
IOPAMIDOL, IOPAMIDOL
LACOSAMIDE, LACOSAMIDE
LEUCOVORIN CALCIUM, LEUCOVORIN CALCIUM
OSELTAMIVIR PHOSPHATE, OSELTAMIVIR PHOSPHATE
VORICONAZOLE, VORICONAZOLE

HAINAN POLY PHARM

* HAINAN POLY PHARMACEUTICAL CO LTD
DAPTOMYCIN, DAPTOMYCIN
GADOBUTROL, GADOBUTROL
LEVETIRACETAM, LEVETIRACETAM
LEVOLEUCOVORIN CALCIUM, LEVOLEUCOVORIN CALCIUM

HALEON US HOLDINGS

* HALEON US HOLDINGS LLC
ABREVA, DOCOSANOL (OTC)
ADVIL ALLERGY AND CONGESTION RELIEF, CHLORPHENIRAMINE MALEATE (OTC)
ADVIL ALLERGY SINUS, CHLORPHENIRAMINE MALEATE (OTC)
ADVIL COLD AND SINUS, IBUPROFEN (OTC)
ADVIL CONGESTION RELIEF, IBUPROFEN (OTC)
ADVIL DUAL ACTION WITH ACETAMINOPHEN, ACETAMINOPHEN (OTC)
ADVIL LIQUI-GELS, IBUPROFEN (OTC)
ADVIL MIGRAINE LIQUI-GELS, IBUPROFEN (OTC)
ADVIL MULTI-SYMPTOM COLD & FLU, CHLORPHENIRAMINE MALEATE (OTC)
ADVIL PM, DIPHENHYDRAMINE CITRATE (OTC)
ADVIL PM, DIPHENHYDRAMINE HYDROCHLORIDE (OTC)
ADVIL, IBUPROFEN (OTC)
ADVIL, IBUPROFEN SODIUM (OTC)
ALLI, ORLISTAT (OTC)
AXID AR, NIZATIDINE (OTC)
CHILDREN'S ADVIL ALLERGY SINUS, CHLORPHENIRAMINE MALEATE (OTC)
CHILDREN'S ADVIL COLD, IBUPROFEN (OTC)
CHILDREN'S ADVIL, IBUPROFEN (OTC)
CHILDREN'S ADVIL-FLAVORED, IBUPROFEN (OTC)
EXCEDRIN (MIGRAINE RELIEF), ACETAMINOPHEN (OTC)
FLONASE ALLERGY RELIEF, FLUTICASONE PROPIONATE (OTC)
FLONASE SENSIMIST ALLERGY RELIEF, FLUTICASONE FUROATE (OTC)
INFANT'S ADVIL, IBUPROFEN (OTC)
JUNIOR STRENGTH ADVIL, IBUPROFEN (OTC)
NICORETTE (MINT), NICOTINE POLACRILEX (OTC)
NICORETTE, NICOTINE POLACRILEX (OTC)
VOLTAREN ARTHRITIS PAIN, DICLOFENAC SODIUM (OTC)

HALOCARBON PRODS

* HALOCARBON PRODUCTS CORP
ISOFLURANE, ISOFLURANE
SEVOFLURANE, SEVOFLURANE

HANGZHOU BINJIANG

* HANGZHOU MINSHENG BINJIANG PHARMACEUTICAL CO LTD
ALENDRONATE SODIUM, ALENDRONATE SODIUM
METHOCARBAMOL, METHOCARBAMOL
TADALAFIL, TADALAFIL

HANGZHOU ZHONGMEI

* HANGZHOU ZHONGMEI HUADONG PHARMACEUTICAL CO LTD
DAPTOMYCIN, DAPTOMYCIN
FONDAPARINUX SODIUM, FONDAPARINUX SODIUM
PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
TACROLIMUS, TACROLIMUS

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** H ******HARM REDUCTION THERP**

* HARM REDUCTION THERAPEUTICS INC
RIVIVE, NALOXONE HYDROCHLORIDE (OTC)

HARMAN FINOCHEM

* HARMAN FINOCHEM LTD
ALLOPURINOL, ALLOPURINOL
BISOPROLOL FUMARATE, BISOPROLOL FUMARATE
METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
THEOPHYLLINE, THEOPHYLLINE

HARMONY

* HARMONY BIOSCIENCES LLC
WAKIX, PITOLISANT HYDROCHLORIDE

HARROW EYE

* HARROW EYE LLC
FLAREX, FLUOROMETHOLONE ACETATE
IHEEZO, CHLOROPROCAINE HYDROCHLORIDE
ILEVRO, NEPAFENAC
IOPIDINE, APRACLONIDINE HYDROCHLORIDE
MAXIDEX, DEXAMETHASONE
MAXITROL, DEXAMETHASONE
NATACYN, NATAMYCIN
NEVANAC, NEPAFENAC
TOBRADEX ST, DEXAMETHASONE
TRIESENCE, TRIAMCINOLONE ACETONIDE
VERKAZIA, CYCLOSPORINE
VEVYE, CYCLOSPORINE
VIGAMOX, MOXIFLOXACIN HYDROCHLORIDE
ZERVIATE, CETIRIZINE HYDROCHLORIDE

HBT LABS INC

* HBT LABS INC
FULVESTRANT, FULVESTRANT

HEC PHARM

* HEC PHARM USA INC
CLARITHROMYCIN, CLARITHROMYCIN
LEVOFLOXACIN, LEVOFLOXACIN
OLANZAPINE, OLANZAPINE
PRASUGREL, PRASUGREL HYDROCHLORIDE

HEC PHARM CO LTD

* HEC PHARM CO LTD
FINGOLIMOD HYDROCHLORIDE, FINGOLIMOD HYDROCHLORIDE

HELSINN

* HELSINN BIREX PHARMACEUTICALS LTD
VALCHLOR, MECHLORETHAMINE HYDROCHLORIDE

HELSINN HLTHCARE

* HELSINN HEALTHCARE SA
AKYNZEO, FOSNETUPITANT CHLORIDE HYDROCHLORIDE
AKYNZEO, NETUPITANT

HERITAGE

* HERITAGE PHARMACEUTICALS INC DBA AVET PHARMACEUTICALS INC
ACYCLOVIR, ACYCLOVIR
AMLODIPINE BESYLATE AND BENAZEPRIL HYDROCHLORIDE, AMLODIPINE BESYLATE
DESIPRAMINE HYDROCHLORIDE, DESIPRAMINE HYDROCHLORIDE
DOCETAXEL, DOCETAXEL
DOXYCYCLINE HCLATE, DOXYCYCLINE HCLATE
DOXYCYCLINE, DOXYCYCLINE
FELODIPINE, FELODIPINE
FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
GLIPIZIDE AND METFORMIN HYDROCHLORIDE, GLIPIZIDE
GLYBURIDE AND METFORMIN HYDROCHLORIDE, GLYBURIDE
GLYBURIDE, GLYBURIDE
HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
LEFLUNOMIDE, LEFLUNOMIDE
NARATRIPTAN, NARATRIPTAN HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** H ****

* HERITAGE PHARMACEUTICALS INC DBA AVET PHARMACEUTICALS INC
 NIMODIPINE, NIMODIPINE
 NYSTATIN, NYSTATIN
 OCTREOTIDE ACETATE, OCTREOTIDE ACETATE
 PAROMOMYCIN SULFATE, PAROMOMYCIN SULFATE
 TORSEMIDE, TORSEMIDE
 TRANEXAMIC ACID, TRANEXAMIC ACID
 VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE
 VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE

HERITAGE LIFE

* HERITAGE LIFE SCIENCES BARBADOS INC
 CLOZARIL, CLOZAPINE

HERITAGE PHARMA

* HERITAGE PHARMA LABS INC
 ACETAMINOPHEN, ACETAMINOPHEN (OTC)
 ACETAZOLAMIDE, ACETAZOLAMIDE
 AMANTADINE HYDROCHLORIDE, AMANTADINE HYDROCHLORIDE
 BENAZEPRIL HYDROCHLORIDE, BENAZEPRIL HYDROCHLORIDE
 BUMETANIDE, BUMETANIDE
 BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
 BUSPIRONE HYDROCHLORIDE, BUSPIRONE HYDROCHLORIDE
 DESMOPRESSIN ACETATE, DESMOPRESSIN ACETATE
 DIFLUNISAL, DIFLUNISAL
 DOXAZOSIN MESYLATE, DOXAZOSIN MESYLATE
 ENALAPRIL MALEATE, ENALAPRIL MALEATE
 HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE
 HYDROXYZINE PAMOATE, HYDROXYZINE PAMOATE
 LABETALOL HYDROCHLORIDE, LABETALOL HYDROCHLORIDE
 LITHIUM CARBONATE, LITHIUM CARBONATE
 METHIMAZOLE, METHIMAZOLE
 NIFEDIPINE, NIFEDIPINE

HERITAGE PHARMA AVET

* HERITAGE PHARMA LABS INC DBA AVET PHARMACEUTICALS LABS INC
 LURASIDONE HYDROCHLORIDE, LURASIDONE HYDROCHLORIDE
 SERTRALINE HYDROCHLORIDE, SERTRALINE HYDROCHLORIDE
 TERAZOSIN HYDROCHLORIDE, TERAZOSIN HYDROCHLORIDE
 THEOPHYLLINE, THEOPHYLLINE
 TRIMETHOBENZAMIDE HYDROCHLORIDE, TRIMETHOBENZAMIDE HYDROCHLORIDE

HERITAGE PHARMS INC

* HERITAGE PHARMACEUTICALS INC
 CALCIUM ACETATE, CALCIUM ACETATE
 ETHOSUXIMIDE, ETHOSUXIMIDE
 HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
 TROSPIMUM CHLORIDE, TROSPIMUM CHLORIDE

HERITAGE PHARMS LABS

* HERITAGE PHARMACEUTICALS LABS INC DBA AVET PHARMACEUTICALS LABS INC
 BENZONATATE, BENZONATATE

HERON THERAPS INC

* HERON THERAPEUTICS INC
 APONVIE, APREPITANT
 CINVANTI, APREPITANT
 SUSTOL, GRANISETRON
 ZYNRELEF KIT, BUPIVACAINE

HETERO LABS

* HETERO LABS LTD
 AMLODIPINE BESYLATE AND VALSARTAN, AMLODIPINE BESYLATE
 DROSPIRENONE AND ETHINYL ESTRADIOL, DROSPIRENONE
 LEVONORGESTREL AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
 LINEZOLID, LINEZOLID
 MELPHALAN HYDROCHLORIDE, MELPHALAN HYDROCHLORIDE

HETERO LABS LTD III

* HETERO LABS LTD UNIT III
 ABACAVIR SULFATE, ABACAVIR SULFATE
 ACYCLOVIR, ACYCLOVIR
 ARIPIPRAZOLE, ARIPIPRAZOLE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** H ****

* HETERO LABS LTD UNIT III
 ATAZANAVIR SULFATE, ATAZANAVIR SULFATE
 ATOVAQUONE, ATOVAQUONE
 CHILDREN'S CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE
 CLOBAZAM, CLOBAZAM
 DABIGATRAN ETEXILATE MESYLATE, DABIGATRAN ETEXILATE MESYLATE
 DARUNAVIR, DARUNAVIR
 DEXTROMETHORPHAN HYDROBROMIDE AND QUINIDINE SULFATE, DEXTROMETHORPHAN HYDROBROMIDE
 DIMETHYL FUMARATE, DIMETHYL FUMARATE
 DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE
 EFAVIRENZ, EFAVIRENZ
 EMTRICITABINE AND TENOFOVIR DISOPROXIL FUMARATE, EMTRICITABINE
 ESCITALOPRAM OXALATE, ESCITALOPRAM OXALATE
 ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM
 ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM (OTC)
 EZETIMIBE, EZETIMIBE
 FENOFIBRATE, FENOFIBRATE
 FINASTERIDE, FINASTERIDE
 HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
 HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE
 INDOMETHACIN, INDOMETHACIN
 KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
 LACOSAMIDE, LACOSAMIDE
 LAMIVUDINE AND ZIDOVUDINE, LAMIVUDINE
 LAMIVUDINE, LAMIVUDINE
 LANSOPRAZOLE, LANSOPRAZOLE
 LEVETIRACETAM, LEVETIRACETAM
 LEVOCETIRIZINE DIHYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE
 LEVOCETIRIZINE DIHYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE (OTC)
 LEVOCETIRIZINE HYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE
 LITHIUM CARBONATE, LITHIUM CARBONATE
 LOPINAVIR AND RITONAVIR, LOPINAVIR
 LORATADINE, LORATADINE (OTC)
 MARAVIROC, MARAVIROC
 METHOCARBAMOL, METHOCARBAMOL
 METOPROLOL SUCCINATE, METOPROLOL SUCCINATE
 NAPROXEN, NAPROXEN
 NEBIVOLOL HYDROCHLORIDE, NEBIVOLOL HYDROCHLORIDE
 NEVIRAPINE, NEVIRAPINE
 OMEPRAZOLE MAGNESIUM, OMEPRAZOLE MAGNESIUM (OTC)
 OMEPRAZOLE, OMEPRAZOLE
 OSELTAMIVIR PHOSPHATE, OSELTAMIVIR PHOSPHATE
 OXCARBAZEPINE, OXCARBAZEPINE
 POSACONAZOLE, POSACONAZOLE
 PREGABALIN, PREGABALIN
 RALTEGRAVIR POTASSIUM, RALTEGRAVIR POTASSIUM
 RANOLAZINE, RANOLAZINE
 RITONAVIR, RITONAVIR
 ROFLUMILAST, ROFLUMILAST
 RUFINAMIDE, RUFINAMIDE
 SIMVASTATIN, SIMVASTATIN
 TADALAFIL, TADALAFIL
 TENOFOVIR DISOPROXIL FUMARATE, TENOFOVIR DISOPROXIL FUMARATE
 TOLTERODINE TARTRATE, TOLTERODINE TARTRATE
 TORSEMIDE, TORSEMIDE
 TRIENTINE HYDROCHLORIDE, TRIENTINE HYDROCHLORIDE
 VARENICLINE TARTRATE, VARENICLINE TARTRATE
 ZIDOVUDINE, ZIDOVUDINE

HETERO LABS LTD V

* HETERO LABS LTD UNIT V
 ACYCLOVIR, ACYCLOVIR
 ALLOPURINOL, ALLOPURINOL
 APIXABAN, APIXABAN
 ARIPIPRAZOLE, ARIPIPRAZOLE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** H ****

* HETERO LABS LTD UNIT V
 ATOMOXETINE HYDROCHLORIDE, ATOMOXETINE HYDROCHLORIDE
 ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM
 AVANAFIL, AVANAFIL
 CAPECITABINE, CAPECITABINE
 CHILDREN'S FEXOFENADINE HYDROCHLORIDE ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
 CINACALCET HYDROCHLORIDE, CINACALCET HYDROCHLORIDE
 CLOPIDOGREL BISULFATE, CLOPIDOGREL BISULFATE
 COLCHICINE, COLCHICINE
 DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
 EFAVIRENZ, EMTRICITABINE, AND TENOFOVIR DISOPROXIL FUMARATE, EFAVIRENZ
 ENTECAVIR, ENTECAVIR
 ERLOTINIB HYDROCHLORIDE, ERLOTINIB HYDROCHLORIDE
 ERTUGLIFLOZIN, ERTUGLIFLOZIN
 ESLICARBAZEPINE ACETATE, ESLICARBAZEPINE ACETATE
 ESZOPICLONE, ESZOPICLONE
 FAMCICLOVIR, FAMCICLOVIR
 FESOTERODINE FUMARATE, FESOTERODINE FUMARATE
 FEXOFENADINE HYDROCHLORIDE ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
 FINGOLIMOD HYDROCHLORIDE, FINGOLIMOD HYDROCHLORIDE
 IRBESARTAN, IRBESARTAN
 LACOSAMIDE, LACOSAMIDE
 LAMIVUDINE AND ZIDOVUDINE, LAMIVUDINE
 LAMIVUDINE, LAMIVUDINE
 LENALIDOMIDE, LENALIDOMIDE
 LEVOFLOXACIN, LEVOFLOXACIN
 LINEZOLID, LINEZOLID
 LOPERAMIDE HYDROCHLORIDE AND SIMETHICONE, LOPERAMIDE HYDROCHLORIDE (OTC)
 LORATADINE, LORATADINE (OTC)
 LOSARTAN POTASSIUM, LOSARTAN POTASSIUM
 MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
 MILNACIPRAN HYDROCHLORIDE, MILNACIPRAN HYDROCHLORIDE
 MONTELUKAST SODIUM, MONTELUKAST SODIUM
 MOXIFLOXACIN HYDROCHLORIDE, MOXIFLOXACIN HYDROCHLORIDE
 MYCOPHENOLATE MOFETIL, MYCOPHENOLATE MOFETIL
 NAPROXEN SODIUM, NAPROXEN SODIUM (OTC)
 OLMESARTAN MEDOXOMIL, OLMESARTAN MEDOXOMIL
 OSELTAMIVIR PHOSPHATE, OSELTAMIVIR PHOSPHATE
 OSPEMIFENE, OSPEMIFENE
 PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
 PIRFENIDONE, PIRFENIDONE
 PITAVASTATIN CALCIUM, PITAVASTATIN CALCIUM
 POMALIDOMIDE, POMALIDOMIDE
 QUETIAPINE FUMARATE, QUETIAPINE FUMARATE
 ROSUVASTATIN CALCIUM, ROSUVASTATIN CALCIUM
 SILDENAFIL CITRATE, SILDENAFIL CITRATE
 SILODOSIN, SILODOSIN
 SIROLIMUS, SIROLIMUS
 SODIUM PICOSULFATE, MAGNESIUM OXIDE AND ANHYDROUS CITRIC ACID, CITRIC ACID
 TELMISARTAN, TELMISARTAN
 TEMOZOLOMIDE, TEMOZOLOMIDE
 TERIFLUNOMIDE, TERIFLUNOMIDE
 TETRABENAZINE, TETRABENAZINE
 TOLTERODINE TARTRATE, TOLTERODINE TARTRATE
 TOLVAPTAN, TOLVAPTAN
 VALACYCLOVIR HYDROCHLORIDE, VALACYCLOVIR HYDROCHLORIDE
 VALGANCICLOVIR HYDROCHLORIDE, VALGANCICLOVIR HYDROCHLORIDE
 VALSARTAN, VALSARTAN

HETERO LABS LTD VI

* HETERO LABS LTD UNIT VI
 AZACITIDINE, AZACITIDINE
 BORTEZOMIB, BORTEZOMIB
 DECITABINE, DECITABINE
 PEMETREXED DISODIUM, PEMETREXED DISODIUM

HEYL CHEMISCH

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** H ****

* HEYL CHEMISCH PHARMAZEUTISCHE FABRIK
RADIOGARDASE (PRUSSIAN BLUE), FERRIC HEXACYANOFERRATE(II)

HIBROW HLTHCARE

* HIBROW HEALTHCARE PRIVATE LTD
ACETAZOLAMIDE, ACETAZOLAMIDE
BACLOFEN, BACLOFEN
HYDROCORTISONE, HYDROCORTISONE
LABETALOL HYDROCHLORIDE, LABETALOL HYDROCHLORIDE
NIACIN, NIACIN
POTASSIUM CITRATE, POTASSIUM CITRATE
QUINIDINE GLUCONATE, QUINIDINE GLUCONATE
URSODIOL, URSODIOL

HIKMA

* HIKMA FARMACEUTICA PORTUGAL SA
BENZTROPINE MESYLATE, BENZTROPINE MESYLATE
BETAMETHASONE ACETATE AND BETAMETHASONE SODIUM PHOSPHATE, BETAMETHASONE ACETATE
CEFTRIAZONE SODIUM, CEFTRIAZONE SODIUM
CEFUROXIME SODIUM, CEFUROXIME SODIUM
DOCETAXEL, DOCETAXEL
ESTRADIOL VALERATE, ESTRADIOL VALERATE
ETOMIDATE, ETOMIDATE
METHYLPREDNISOLONE SODIUM SUCCINATE, METHYLPREDNISOLONE SODIUM SUCCINATE
NOREPINEPHRINE BITARTRATE, NOREPINEPHRINE BITARTRATE
SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE
VECURONIUM BROMIDE, VECURONIUM BROMIDE

* HIKMA PHARMACEUTICALS
AMOXICILLIN, AMOXICILLIN
CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE

* HIKMA PHARMACEUTICALS INTERNATIONAL LTD
CODEINE SULFATE, CODEINE SULFATE
DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE
DIGOXIN, DIGOXIN
FUROSEMIDE, FUROSEMIDE
GEMCITABINE HYDROCHLORIDE, GEMCITABINE HYDROCHLORIDE
MICAfungin SODIUM, MICAfungin SODIUM
MIDAZOLAM HYDROCHLORIDE, MIDAZOLAM HYDROCHLORIDE
MORPHINE SULFATE, MORPHINE SULFATE
RUFINAMIDE, RUFINAMIDE
STERILE WATER FOR INJECTION, STERILE WATER FOR INJECTION
SUCCINYLCHOLINE CHLORIDE, SUCCINYLCHOLINE CHLORIDE

* HIKMA PHARMACEUTICALS LLC
ABIRATERONE ACETATE, ABIRATERONE ACETATE
DANTROLENE SODIUM, DANTROLENE SODIUM
DOXERCALCIFEROL, DOXERCALCIFEROL
PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM

* HIKMA PHARMACEUTICALS USA INC
ACARBOSE, ACARBOSE
ACETAMINOPHEN, ACETAMINOPHEN
ACETAZOLAMIDE SODIUM, ACETAZOLAMIDE SODIUM
ADENOSINE, ADENOSINE
ALENDRONATE SODIUM, ALENDRONATE SODIUM
ALLOPURINOL SODIUM, ALLOPURINOL SODIUM
ALPRAZOLAM, ALPRAZOLAM
ALPROSTADIL, ALPROSTADIL
ALVIMOPAN, ALVIMOPAN
AMICAR, AMINOCAPROIC ACID
AMIKACIN SULFATE, AMIKACIN SULFATE
AMPICILLIN AND SULBACTAM, AMPICILLIN SODIUM
AMRINONE LACTATE, INAMRINONE LACTATE
ATIVAN, LORAZEPAM
ATRACURIUM BESYLATE PRESERVATIVE FREE, ATRACURIUM BESYLATE
ATRACURIUM BESYLATE, ATRACURIUM BESYLATE
ATROPINE SULFATE, ATROPINE SULFATE
AZATHIOPRINE SODIUM, AZATHIOPRINE SODIUM

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** H ****

* HIKMA PHARMACEUTICALS USA INC
 BALSALAZIDE DISODIUM, BALSALAZIDE DISODIUM
 BEXAROTENE, BEXAROTENE
 BLEOMYCIN SULFATE, BLEOMYCIN SULFATE
 BUPRENORPHINE HYDROCHLORIDE AND NALOXONE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE
 BUPRENORPHINE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE
 BUTALBITAL, ACETAMINOPHEN, CAFFEINE AND CODEINE PHOSPHATE, ACETAMINOPHEN
 BUTORPHANOL TARTRATE PRESERVATIVE FREE, BUTORPHANOL TARTRATE
 BUTORPHANOL TARTRATE, BUTORPHANOL TARTRATE
 CAFKIT, CAFFEINE CITRATE
 CALCITRIOL, CALCITRIOL
 CALCIUM ACETATE, CALCIUM ACETATE
 CARBOPLATIN, CARBOPLATIN
 CEFAZOLIN SODIUM, CEFAZOLIN SODIUM
 CEFEPIME HYDROCHLORIDE, CEFEPIME HYDROCHLORIDE
 CEFOXITIN, CEFOXITIN SODIUM
 CERUBIDINE, DAUNORUBICIN HYDROCHLORIDE
 CEVIMELINE HYDROCHLORIDE, CEVIMELINE HYDROCHLORIDE
 CHLOROPROCAINE HYDROCHLORIDE, CHLOROPROCAINE HYDROCHLORIDE
 CHLORPROMAZINE HYDROCHLORIDE, CHLORPROMAZINE HYDROCHLORIDE
 CISATRACURIUM BESYLATE, CISATRACURIUM BESYLATE
 CISPLATIN, CISPLATIN
 CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE
 CLADRIBINE, CLADRIBINE
 CLINDAMYCIN PHOSPHATE IN DEXTROSE 5%, CLINDAMYCIN PHOSPHATE
 CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
 CLOTRIMAZOLE, CLOTRIMAZOLE
 COMBOGESIC IV, ACETAMINOPHEN
 CYCLOPHOSPHAMIDE, CYCLOPHOSPHAMIDE
 CYCLOSPORINE, CYCLOSPORINE
 CYTARABINE, CYTARABINE
 DACARBAZINE, DACARBAZINE
 DAPTOMYCIN, DAPTOMYCIN
 DAUNORUBICIN HYDROCHLORIDE, DAUNORUBICIN HYDROCHLORIDE
 DEFERIPRONE, DEFERIPRONE
 DEFEROXAMINE MESYLATE, DEFEROXAMINE MESYLATE
 DESVENLAFAXINE SUCCINATE, DESVENLAFAXINE SUCCINATE
 DEXAMETHASONE INTENSOL, DEXAMETHASONE
 DEXAMETHASONE SODIUM PHOSPHATE, DEXAMETHASONE SODIUM PHOSPHATE
 DEXAMETHASONE, DEXAMETHASONE
 DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE
 DEXRAZOXANE HYDROCHLORIDE, DEXRAZOXANE HYDROCHLORIDE
 DIAZEPAM INTENSOL, DIAZEPAM
 DIAZEPAM, DIAZEPAM
 DICYCLOMINE HYDROCHLORIDE (PRESERVATIVE FREE), DICYCLOMINE HYDROCHLORIDE
 DICYCLOMINE HYDROCHLORIDE, DICYCLOMINE HYDROCHLORIDE
 DIGOXIN, DIGOXIN
 DIHYDROERGOTAMINE MESYLATE, DIHYDROERGOTAMINE MESYLATE
 DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
 DIPHENOXYLATE HYDROCHLORIDE AND ATROPINE SULFATE, ATROPINE SULFATE
 DIPYRIDAMOLE, DIPYRIDAMOLE
 DOBUTAMINE HYDROCHLORIDE, DOBUTAMINE HYDROCHLORIDE
 DOPRAM, DOXAPRAM HYDROCHLORIDE
 DOXORUBICIN HYDROCHLORIDE, DOXORUBICIN HYDROCHLORIDE
 DROPERIDOL, DROPERIDOL
 DURAMORPH PF, MORPHINE SULFATE
 EDARAVONE, EDARAVONE
 EPHEDRINE SULFATE, EPHEDRINE SULFATE
 EPIRUBICIN HYDROCHLORIDE, EPIRUBICIN HYDROCHLORIDE
 ESMOLOL HYDROCHLORIDE, ESMOLOL HYDROCHLORIDE
 ETOMIDATE, ETOMIDATE
 ETOPOSIDE, ETOPOSIDE
 EVEROLIMUS, EVEROLIMUS
 EXEMESTANE, EXEMESTANE
 FAMOTIDINE PRESERVATIVE FREE, FAMOTIDINE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** H ****

* HIKMA PHARMACEUTICALS USA INC
 FAMOTIDINE, FAMOTIDINE
 FEBUXOSTAT, FEBUXOSTAT
 FENTANYL CITRATE, FENTANYL CITRATE
 FLECAINIDE ACETATE, FLECAINIDE ACETATE
 FLOXURIDINE, FLOXURIDINE
 FLUCONAZOLE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, FLUCONAZOLE
 FLUCONAZOLE IN SODIUM CHLORIDE 0.9%, FLUCONAZOLE
 FLUDARABINE PHOSPHATE, FLUDARABINE PHOSPHATE
 FLUMAZENIL, FLUMAZENIL
 FLUPHENAZINE DECANOATE, FLUPHENAZINE DECANOATE
 FLUTICASONE PROPIONATE AND SALMETEROL XINAFOATE, FLUTICASONE PROPIONATE
 FLUTICASONE PROPIONATE, FLUTICASONE PROPIONATE
 FLUTICASONE PROPIONATE, FLUTICASONE PROPIONATE (OTC)
 FOSPHENYTOIN SODIUM, FOSPHENYTOIN SODIUM
 FUROSEMIDE, FUROSEMIDE
 GALANTAMINE HYDROBROMIDE, GALANTAMINE HYDROBROMIDE
 GANCICLOVIR SODIUM, GANCICLOVIR SODIUM
 GENTAMICIN SULFATE, GENTAMICIN SULFATE
 GRANISETRON HYDROCHLORIDE, GRANISETRON HYDROCHLORIDE
 HALOPERIDOL DECANOATE, HALOPERIDOL DECANOATE
 HALOPERIDOL, HALOPERIDOL LACTATE
 HEPARIN SODIUM, HEPARIN SODIUM
 HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
 HYDROMORPHONE HYDROCHLORIDE, HYDROMORPHONE HYDROCHLORIDE
 ICOSAPENT ETHYL, ICOSAPENT ETHYL
 IDARUBICIN HYDROCHLORIDE, IDARUBICIN HYDROCHLORIDE
 IFOSFAMIDE, IFOSFAMIDE
 IMIPRAMINE PAMOATE, IMIPRAMINE PAMOATE
 IMMPHENTIV, PHENYLEPHRINE HYDROCHLORIDE
 INDOMETHACIN, INDOMETHACIN
 INFUMORPH, MORPHINE SULFATE
 IPRATROPIUM BROMIDE, IPRATROPIUM BROMIDE
 IRBESARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE
 ISOPROTERENOL HYDROCHLORIDE, ISOPROTERENOL HYDROCHLORIDE
 KETAMINE HYDROCHLORIDE, KETAMINE HYDROCHLORIDE
 KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
 KLOXXADO, NALOXONE HYDROCHLORIDE
 LABETALOL HYDROCHLORIDE IN DEXTROSE, LABETALOL HYDROCHLORIDE
 LABETALOL HYDROCHLORIDE IN SODIUM CHLORIDE, LABETALOL HYDROCHLORIDE
 LABETALOL HYDROCHLORIDE, LABETALOL HYDROCHLORIDE
 LACOSAMIDE, LACOSAMIDE
 LEUCOVORIN CALCIUM PRESERVATIVE FREE, LEUCOVORIN CALCIUM
 LEUCOVORIN CALCIUM, LEUCOVORIN CALCIUM
 LEVETIRACETAM IN SODIUM CHLORIDE, LEVETIRACETAM
 LEVOCARNITINE, LEVOCARNITINE
 LEVORPHANOL TARTRATE, LEVORPHANOL TARTRATE
 LEVOTHYROXINE SODIUM, LEVOTHYROXINE SODIUM
 LIDOCAINE HYDROCHLORIDE PRESERVATIVE FREE, LIDOCAINE HYDROCHLORIDE
 LIDOCAINE HYDROCHLORIDE, LIDOCAINE HYDROCHLORIDE
 LIDOCAINE VISCOUS, LIDOCAINE HYDROCHLORIDE
 LINEZOLID, LINEZOLID
 LIRAGLUTIDE, LIRAGLUTIDE
 LISDEXAMFETAMINE DIMESYLATE, LISDEXAMFETAMINE DIMESYLATE
 LITHIUM CARBONATE, LITHIUM CARBONATE
 LORAZEPAM INTENSOL, LORAZEPAM
 MEFLOQUINE HYDROCHLORIDE, MEFLOQUINE HYDROCHLORIDE
 MELPHALAN HYDROCHLORIDE, MELPHALAN HYDROCHLORIDE
 MEPERIDINE HYDROCHLORIDE, MEPERIDINE HYDROCHLORIDE
 MERCAPTOPYRINE, MERCAPTOPYRINE
 MESNA, MESNA
 METHADONE HYDROCHLORIDE INTENSOL, METHADONE HYDROCHLORIDE
 METHADONE HYDROCHLORIDE, METHADONE HYDROCHLORIDE
 METHAMPHETAMINE HYDROCHLORIDE, METHAMPHETAMINE HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** H ****

* HIKMA PHARMACEUTICALS USA INC
 METHOTREXATE SODIUM PRESERVATIVE FREE, METHOTREXATE SODIUM
 METHOTREXATE SODIUM, METHOTREXATE SODIUM
 METOPROLOL TARTRATE, METOPROLOL TARTRATE
 MIDAZOLAM HYDROCHLORIDE, MIDAZOLAM HYDROCHLORIDE
 MIDAZOLAM IN 0.9% SODIUM CHLORIDE, MIDAZOLAM
 MILRINONE LACTATE IN DEXTROSE 5% IN PLASTIC CONTAINER, MILRINONE LACTATE
 MILRINONE LACTATE, MILRINONE LACTATE
 MITOMYCIN, MITOMYCIN
 MITOXANTRONE HYDROCHLORIDE, MITOXANTRONE HYDROCHLORIDE
 MORPHINE SULFATE, MORPHINE SULFATE
 MYCOPHENOLATE MOFETIL, MYCOPHENOLATE MOFETIL
 NALOXONE HYDROCHLORIDE, NALOXONE HYDROCHLORIDE
 NALOXONE, NALOXONE HYDROCHLORIDE
 NAPROXEN, NAPROXEN
 NARATRIPTAN, NARATRIPTAN HYDROCHLORIDE
 NEOSTIGMINE METHYLSULFATE, NEOSTIGMINE METHYLSULFATE
 NOREPINEPHRINE BITARTRATE, NOREPINEPHRINE BITARTRATE
 ONDANSETRON HYDROCHLORIDE PRESERVATIVE FREE, ONDANSETRON HYDROCHLORIDE
 ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
 ORPHENADRINE CITRATE, ORPHENADRINE CITRATE
 OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
 OXYMORPHONE HYDROCHLORIDE, OXYMORPHONE HYDROCHLORIDE
 OXYTOCIN, OXYTOCIN
 PACLITAXEL, PACLITAXEL
 PAMIDRONATE DISODIUM, PAMIDRONATE DISODIUM
 PENTOBARBITAL SODIUM, PENTOBARBITAL SODIUM
 PHENTOLAMINE MESYLATE, PHENTOLAMINE MESYLATE
 PHENYLEPHRINE HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE
 PHENYTOIN SODIUM, PHENYTOIN SODIUM
 PHYTONADIONE, PHYTONADIONE
 POSACONAZOLE, POSACONAZOLE
 PREDNISONE INTENSOL, PREDNISONE
 PREDNISONE, PREDNISONE
 PROCHLORPERAZINE EDISYLATE, PROCHLORPERAZINE EDISYLATE
 PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
 PROPOFOL, PROPOFOL
 PROPRANOLOL HYDROCHLORIDE, PROPRANOLOL HYDROCHLORIDE
 PROTRIPTYLINE HYDROCHLORIDE, PROTRIPTYLINE HYDROCHLORIDE
 PYRAZINAMIDE, PYRAZINAMIDE
 QUETIAPINE FUMARATE, QUETIAPINE FUMARATE
 RAMIPRIL, RAMIPRIL
 REGADENOSON, REGADENOSON
 REMIFENTANIL HYDROCHLORIDE, REMIFENTANIL HYDROCHLORIDE
 RIFAMPIN, RIFAMPIN
 ROBAXIN, METHOCARBAMOL
 ROPIVACAINE HYDROCHLORIDE, ROPIVACAINE HYDROCHLORIDE
 RUFINAMIDE, RUFINAMIDE
 SODIUM ACETATE, SODIUM ACETATE
 SODIUM CHLORIDE 0.9%, SODIUM CHLORIDE
 SODIUM FERRIC GLUCONATE COMPLEX IN SUCROSE, FERRIC OXYHYDROXIDE
 SODIUM TETRADECYL SULFATE, SODIUM TETRADECYL SULFATE
 STERILE WATER FOR INJECTION, STERILE WATER FOR INJECTION
 SUCCINYLCHOLINE CHLORIDE, SUCCINYLCHOLINE CHLORIDE
 SUFENTANIL CITRATE, SUFENTANIL CITRATE
 SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
 TESTOSTERONE CYPIONATE, TESTOSTERONE CYPIONATE
 THIOTEPA, THIOTEPA
 TOBRAMYCIN SULFATE, TOBRAMYCIN SULFATE
 TOBRAMYCIN, TOBRAMYCIN
 TORSEMIDE, TORSEMIDE
 TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE
 VALRUBICIN, VALRUBICIN
 VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE
 VECURONIUM BROMIDE, VECURONIUM BROMIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** H ****

* HIKMA PHARMACEUTICALS USA INC
 VIGABATRIN, VIGABATRIN
 VINBLASTINE SULFATE, VINBLASTINE SULFATE
 VINOURELBINE TARTRATE, VINOURELBINE TARTRATE
 VORICONAZOLE, VORICONAZOLE
 ZALEPLON, ZALEPLON

HIKMA FARMACEUTICA

* HIKMA FARMACEUTICA (PORTUGAL) SA
 AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE
 BENZTROPINE MESYLATE, BENZTROPINE MESYLATE
 BUTORPHANOL TARTRATE, BUTORPHANOL TARTRATE
 CEFOXITIN, CEFOXITIN SODIUM
 CEFTRIAZONE, CEFTRIAZONE SODIUM
 CIPROFLOXACIN IN DEXTROSE 5% IN PLASTIC CONTAINER, CIPROFLOXACIN
 CIPROFLOXACIN, CIPROFLOXACIN
 CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
 ENALAPRILAT, ENALAPRILAT
 FLUCONAZOLE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, FLUCONAZOLE
 FLUMAZENIL, FLUMAZENIL
 FOSPHENYTOIN SODIUM, FOSPHENYTOIN SODIUM
 GLYCOPYRROLATE, GLYCOPYRROLATE
 GRANISETRON HYDROCHLORIDE, GRANISETRON HYDROCHLORIDE
 IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE
 LEVETIRACETAM, LEVETIRACETAM
 LEVOFLOXACIN IN DEXTROSE 5% IN PLASTIC CONTAINER, LEVOFLOXACIN
 METOPROLOL TARTRATE, METOPROLOL TARTRATE
 MILRINONE LACTATE IN PLASTIC CONTAINER, MILRINONE LACTATE
 MILRINONE LACTATE, MILRINONE LACTATE
 ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
 PROGESTERONE, PROGESTERONE
 PROPRANOLOL HYDROCHLORIDE, PROPRANOLOL HYDROCHLORIDE
 TERBUTALINE SULFATE, TERBUTALINE SULFATE
 TESTOSTERONE CYPIONATE, TESTOSTERONE CYPIONATE
 VALPROATE SODIUM, VALPROATE SODIUM

* HIKMA FARMACEUTICA PORTUGAL LDA
 CEFAZOLIN SODIUM, CEFAZOLIN SODIUM
 FLUCONAZOLE IN SODIUM CHLORIDE 0.9%, FLUCONAZOLE

* HIKMA FARMACEUTICA PORTUGAL SA
 OXYTOCIN, OXYTOCIN
 TESTOSTERONE ENANTHATE, TESTOSTERONE ENANTHATE

* HIKMA FARMACEUTICA SA
 ZOLEDRONIC ACID, ZOLEDRONIC ACID

HIKMA INTL PHARMS

* HIKMA INTERNATIONAL PHARMACEUTICALS LLC
 CAPTOPRIL, CAPTOPRIL
 DIGOXIN, DIGOXIN
 DOPAMINE HYDROCHLORIDE, DOPAMINE HYDROCHLORIDE
 DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
 GEMCITABINE HYDROCHLORIDE, GEMCITABINE HYDROCHLORIDE
 ISOSORBIDE DINITRATE, ISOSORBIDE DINITRATE
 MITIGARE, COLCHICINE
 NICARDIPINE HYDROCHLORIDE IN 0.9% SODIUM CHLORIDE, NICARDIPINE HYDROCHLORIDE
 NICARDIPINE HYDROCHLORIDE, NICARDIPINE HYDROCHLORIDE

HIKMA PHARM CO LTD

* HIKMA PHARM CO LTD
 ARGATROBAN, ARGATROBAN

HIKMA PHARMS

* HIKMA PHARMACEUTICALS
 AMOXICILLIN AND CLAVULANATE POTASSIUM, AMOXICILLIN
 AMOXICILLIN, AMOXICILLIN
 BUPIVACAINE HYDROCHLORIDE PRESERVATIVE FREE, BUPIVACAINE HYDROCHLORIDE
 BUPIVACAINE HYDROCHLORIDE, BUPIVACAINE HYDROCHLORIDE
 DICYCLOMINE HYDROCHLORIDE, DICYCLOMINE HYDROCHLORIDE
 DIHYDROERGOTAMINE MESYLATE, DIHYDROERGOTAMINE MESYLATE
 PENICILLIN V POTASSIUM, PENICILLIN V POTASSIUM

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** H ****

- * HIKMA PHARMACEUTICALS
RIFAMPIN, RIFAMPIN
VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE
- * HIKMA PHARMACEUTICALS CO LTD
PARICALCITOL, PARICALCITOL
- HILL DERMAC**
- * HILL DERMACEUTICALS INC
DERMA-SMOOTH/FS, FLUOCINOLONE ACETONIDE
DERMOTIC, FLUOCINOLONE ACETONIDE
- HILL DERMACEUTICALS**
- * HILL DERMACEUTICALS INC
TOLAK, FLUOROURACIL
- HISAMITSU**
- * HISAMITSU PHARMACEUTICAL CO INC
SECUADO, ASENAPINE
- HISAMITSU PHARM CO**
- * HISAMITSU PHARMACEUTICAL CO INC
SALONPAS, MENTHOL (OTC)
- HISUN PHARM HANGZHOU**
- * HISUN PHARMACEUTICAL (HANGZHOU) CO LTD
CLADRIBINE, CLADRIBINE
EPIRUBICIN HYDROCHLORIDE, EPIRUBICIN HYDROCHLORIDE
IRBESARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
IRBESARTAN, IRBESARTAN
LEVETIRACETAM, LEVETIRACETAM
OLANZAPINE, OLANZAPINE
- * HISUN PHARMACEUTICAL HANGZHOU CO LTD
DACTINOMYCIN, DACTINOMYCIN
DAPTOMYCIN, DAPTOMYCIN
DAUNORUBICIN HYDROCHLORIDE, DAUNORUBICIN HYDROCHLORIDE
DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
TICAGRELOR, TICAGRELOR
- HLTHCARE**
- * HEALTHCARE PHARMACEUTICALS LTD
ATENOLOL, ATENOLOL
DROSPIRENONE AND ETHINYL ESTRADIOL, DROSPIRENONE
METHOCARBAMOL, METHOCARBAMOL
- HOFFMANN LA ROCHE**
- * HOFFMANN LA ROCHE INC
ZELBORAF, VEMURAFENIB
- HOFFMANN-LA ROCHE**
- * HOFFMANN-LA ROCHE INC
ALECENSA, ALECTINIB HYDROCHLORIDE
- HONG KONG**
- * HONG KONG KING FRIEND INDUSTRIAL CO LTD
EPTIFIBATIDE, EPTIFIBATIDE
MEDROXYPROGESTERONE ACETATE, MEDROXYPROGESTERONE ACETATE
METHYLPREDNISOLONE ACETATE, METHYLPREDNISOLONE ACETATE
XENLETA, LEFAMULIN ACETATE
- HOPE PHARMS**
- * HOPE PHARMACEUTICALS
NITHIODE, SODIUM NITRITE
SODIUM NITRITE, SODIUM NITRITE
SODIUM THIOSULFATE, SODIUM THIOSULFATE
- HORIZON**
- * HORIZON THERAPEUTICS USA INC
PROCYSBI, CYSTEAMINE BITARTRATE
- HORIZON THERAP US**
- * HORIZON THERAPEUTICS US HOLDING LLC
BUPHENYL, SODIUM PHENYLBUTYRATE
RAVICTI, GLYCEROL PHENYLBUTYRATE
- HOSPIRA**
- * HOSPIRA INC
ACETYLCYSTEINE, ACETYLCYSTEINE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** H ******* HOSPIRA INC**

ALFENTANIL, ALFENTANIL HYDROCHLORIDE
 AMIDATE, ETOMIDATE
 AMINOCAPROIC ACID IN PLASTIC CONTAINER, AMINOCAPROIC ACID
 AMINOPHYLLINE, AMINOPHYLLINE
 AMMONIUM CHLORIDE IN PLASTIC CONTAINER, AMMONIUM CHLORIDE
 ARGATROBAN, ARGATROBAN
 ATROPINE SULFATE, ATROPINE SULFATE
 AZITHROMYCIN, AZITHROMYCIN
 AZTREONAM, AZTREONAM
 BACTERIOSTATIC SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, SODIUM CHLORIDE
 BACTERIOSTATIC WATER FOR INJECTION IN PLASTIC CONTAINER, STERILE WATER FOR INJECTION
 BLEOMYCIN SULFATE, BLEOMYCIN SULFATE
 BORTEZOMIB, BORTEZOMIB
 BUPIVACAINE HYDROCHLORIDE AND EPINEPHRINE, BUPIVACAINE HYDROCHLORIDE
 BUPIVACAINE HYDROCHLORIDE, BUPIVACAINE HYDROCHLORIDE
 BUPRENORPHINE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE
 BUSULFAN, BUSULFAN
 BUTORPHANOL TARTRATE PRESERVATIVE FREE, BUTORPHANOL TARTRATE
 CALCIUM CHLORIDE 10% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 CARBOCAINE, MEPIVACAINE HYDROCHLORIDE
 CARBOPLATIN, CARBOPLATIN
 CHROMIC CHLORIDE IN PLASTIC CONTAINER, CHROMIC CHLORIDE
 CIPROFLOXACIN IN DEXTROSE 5% IN PLASTIC CONTAINER, CIPROFLOXACIN
 CISATRACURIUM BESYLATE, CISATRACURIUM BESYLATE
 CUPRIC CHLORIDE IN PLASTIC CONTAINER, CUPRIC CHLORIDE
 CYTARABINE, CYTARABINE
 DACARBAZINE, DACARBAZINE
 DAPTOMYCIN, DAPTOMYCIN
 DEFEROXAMINE MESYLATE, DEFEROXAMINE MESYLATE
 DEMEROL, MEPERIDINE HYDROCHLORIDE
 DEXTROSE 25%, DEXTROSE
 DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 50%, DEXTROSE
 DIAZEPAM, DIAZEPAM
 DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
 DOBUTAMINE HYDROCHLORIDE IN DEXTROSE 5% IN PLASTIC CONTAINER, DOBUTAMINE HYDROCHLORIDE
 DOBUTAMINE HYDROCHLORIDE, DOBUTAMINE HYDROCHLORIDE
 DOPAMINE HYDROCHLORIDE IN DEXTROSE 5% IN PLASTIC CONTAINER, DOPAMINE HYDROCHLORIDE
 DOPAMINE HYDROCHLORIDE, DOPAMINE HYDROCHLORIDE
 DOXERCALCIFEROL, DOXERCALCIFEROL
 ENALAPRILAT, ENALAPRILAT
 EPINEPHRINE, EPINEPHRINE
 ERYTHROCIN, ERYTHROMYCIN LACTOBIONATE
 FENTANYL CITRATE, FENTANYL CITRATE
 FLUCONAZOLE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, FLUCONAZOLE
 FUROSEMIDE, FUROSEMIDE
 GENTAMICIN SULFATE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, GENTAMICIN SULFATE
 GENTAMICIN SULFATE, GENTAMICIN SULFATE
 HEPARIN SODIUM 1,000 UNITS IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, HEPARIN
 HEPARIN SODIUM 10,000 UNITS IN DEXTROSE 5% IN PLASTIC CONTAINER, HEPARIN SODIUM
 HEPARIN SODIUM 12,500 UNITS IN DEXTROSE 5% IN PLASTIC CONTAINER, HEPARIN SODIUM
 HEPARIN SODIUM 12,500 UNITS IN SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, HEPARIN
 HEPARIN SODIUM 2,000 UNITS IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, HEPARIN
 HEPARIN SODIUM 25,000 UNITS IN DEXTROSE 5% IN PLASTIC CONTAINER, HEPARIN SODIUM
 HEPARIN SODIUM 25,000 UNITS IN SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, HEPARIN
 HEPARIN SODIUM PRESERVATIVE FREE, HEPARIN SODIUM
 HEPARIN SODIUM, HEPARIN SODIUM
 IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE
 KETAMINE HYDROCHLORIDE, KETAMINE HYDROCHLORIDE
 KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
 LABETALOL HYDROCHLORIDE, LABETALOL HYDROCHLORIDE
 LEVOFLOXACIN IN DEXTROSE 5% IN PLASTIC CONTAINER, LEVOFLOXACIN
 LEVOPHED, NOREPINEPHRINE BITARTRATE
 LIDOCAINE HYDROCHLORIDE 5% AND DEXTROSE 7.5%, LIDOCAINE HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** H **

* HOSPIRA INC

LIDOCAINE HYDROCHLORIDE AND EPINEPHRINE, EPINEPHRINE
 LIDOCAINE HYDROCHLORIDE IN PLASTIC CONTAINER, LIDOCAINE HYDROCHLORIDE
 LIDOCAINE HYDROCHLORIDE PRESERVATIVE FREE IN PLASTIC CONTAINER, LIDOCAINE
 LIDOCAINE HYDROCHLORIDE PRESERVATIVE FREE, LIDOCAINE HYDROCHLORIDE
 LIDOCAINE HYDROCHLORIDE, LIDOCAINE HYDROCHLORIDE
 LINEZOLID IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, LINEZOLID
 LORAZEPAM, LORAZEPAM
 MAGNESIUM SULFATE IN DEXTROSE 5% IN PLASTIC CONTAINER, MAGNESIUM SULFATE
 MAGNESIUM SULFATE IN PLASTIC CONTAINER, MAGNESIUM SULFATE
 MAGNESIUM SULFATE, MAGNESIUM SULFATE
 MANGANESE CHLORIDE IN PLASTIC CONTAINER, MANGANESE CHLORIDE
 MANNITOL 25%, MANNITOL
 MARCAINE HYDROCHLORIDE PRESERVATIVE FREE, BUPIVACAINE HYDROCHLORIDE
 MARCAINE HYDROCHLORIDE W/ EPINEPHRINE PRESERVATIVE FREE, BUPIVACAINE HYDROCHLORIDE
 MARCAINE HYDROCHLORIDE W/ EPINEPHRINE, BUPIVACAINE HYDROCHLORIDE
 MARCAINE HYDROCHLORIDE, BUPIVACAINE HYDROCHLORIDE
 MARCAINE, BUPIVACAINE HYDROCHLORIDE
 METHOTREXATE SODIUM PRESERVATIVE FREE, METHOTREXATE SODIUM
 METHOTREXATE SODIUM, METHOTREXATE SODIUM
 METOCLOPRAMIDE HYDROCHLORIDE, METOCLOPRAMIDE HYDROCHLORIDE
 METOPROLOL TARTRATE, METOPROLOL TARTRATE
 METRONIDAZOLE IN PLASTIC CONTAINER, METRONIDAZOLE
 MIDAZOLAM HYDROCHLORIDE PRESERVATIVE FREE, MIDAZOLAM HYDROCHLORIDE
 MIDAZOLAM HYDROCHLORIDE, MIDAZOLAM HYDROCHLORIDE
 MILRINONE LACTATE, MILRINONE LACTATE
 MORPHINE SULFATE, MORPHINE SULFATE
 NALBUPHINE HYDROCHLORIDE, NALBUPHINE HYDROCHLORIDE
 NALOXONE HYDROCHLORIDE, NALOXONE HYDROCHLORIDE
 ONDANSETRON HYDROCHLORIDE PRESERVATIVE FREE, ONDANSETRON HYDROCHLORIDE
 ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
 PACLITAXEL, PACLITAXEL
 PALONOSETRON HYDROCHLORIDE, PALONOSETRON HYDROCHLORIDE
 PAMIDRONATE DISODIUM, PAMIDRONATE DISODIUM
 PEMETREXED DISODIUM, PEMETREXED DISODIUM
 PEMETREXED DITROMETHAMINE, PEMETREXED DITROMETHAMINE
 PLEGISOL IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 POTASSIUM ACETATE, POTASSIUM ACETATE
 POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
 PRECEDEX, DEXMEDETOMIDINE HYDROCHLORIDE
 PROCAINAMIDE HYDROCHLORIDE, PROCAINAMIDE HYDROCHLORIDE
 PROPOFOL, PROPOFOL
 QUELICIN, SUCCINYLCHOLINE CHLORIDE
 REGADENOSON, REGADENOSON
 ROCURONIUM BROMIDE, ROCURONIUM BROMIDE
 SODIUM ACETATE, SODIUM ACETATE
 SODIUM BICARBONATE, SODIUM BICARBONATE
 SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, SODIUM CHLORIDE
 SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, SODIUM CHLORIDE
 SODIUM CHLORIDE 14.6%, SODIUM CHLORIDE
 SODIUM CHLORIDE 23.4%, SODIUM CHLORIDE
 SODIUM PHOSPHATES IN PLASTIC CONTAINER, SODIUM PHOSPHATE, DIBASIC, HEPTAHYDRATE
 STERILE WATER FOR INJECTION, STERILE WATER FOR INJECTION
 SUFENTANIL CITRATE, SUFENTANIL CITRATE
 TAZICEF, CEFTAZIDIME
 THAM, TROMETHAMINE
 TOBRAMYCIN SULFATE, TOBRAMYCIN SULFATE
 TPN ELECTROLYTES IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE
 VECURONIUM BROMIDE, VECURONIUM BROMIDE
 VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE
 VITAMIN K1, PHYTONADIONE
 ZINC CHLORIDE IN PLASTIC CONTAINER, ZINC CHLORIDE
 ZOLEDRONIC ACID, ZOLEDRONIC ACID

* HOSPIRA WORLDWIDE, INC

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** H ****

* HOSPIRA WORLDWIDE, INC
 GEMCITABINE HYDROCHLORIDE, GEMCITABINE HYDROCHLORIDE
 MITOXANTRONE HYDROCHLORIDE, MITOXANTRONE HYDROCHLORIDE
 VINCRISTINE SULFATE PFS, VINCRISTINE SULFATE

HOSPIRA INC

* HOSPIRA INC
 ATRACURIUM BESYLATE PRESERVATIVE FREE, ATRACURIUM BESYLATE
 ATRACURIUM BESYLATE, ATRACURIUM BESYLATE
 DOCETAXEL, DOCETAXEL
 GEMCITABINE HYDROCHLORIDE, GEMCITABINE HYDROCHLORIDE
 HEPARIN SODIUM, HEPARIN SODIUM
 HYDROMORPHONE HYDROCHLORIDE, HYDROMORPHONE HYDROCHLORIDE
 LEVETIRACETAM, LEVETIRACETAM
 MORPHINE SULFATE, MORPHINE SULFATE
 NIPENT, PENTOSTATIN
 TOPOTECAN HYDROCHLORIDE, TOPOTECAN HYDROCHLORIDE
 VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE
 ZOLEDRONIC ACID, ZOLEDRONIC ACID

HOSPIRA WORLDWIDE

* HOSPIRA WORLDWIDE PTY
 OXALIPLATIN, OXALIPLATIN

HOT SHOTS NM LLC

* HOT SHOTS NUCLEAR MEDICINE LLC
 SODIUM FLUORIDE F-18, SODIUM FLUORIDE F-18

HQ SPCLT PHARMA

* HQ SPECIALTY PHARMA CORP
 CALCIUM GLUCONATE IN SODIUM CHLORIDE, CALCIUM GLUCONATE
 CEFAZOLIN SODIUM, CEFAZOLIN SODIUM
 CISPLATIN, CISPLATIN
 DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE
 ESMOLOL HYDROCHLORIDE DOUBLE STRENGTH IN PLASTIC CONTAINER, ESMOLOL HYDROCHLORIDE
 ESMOLOL HYDROCHLORIDE IN PLASTIC CONTAINER, ESMOLOL HYDROCHLORIDE
 ESMOLOL HYDROCHLORIDE, ESMOLOL HYDROCHLORIDE
 IMIPENEM AND CILASTATIN, CILASTATIN SODIUM
 LEVETIRACETAM IN SODIUM CHLORIDE, LEVETIRACETAM
 LINEZOLID, LINEZOLID
 MAGNESIUM SULFATE IN DEXTROSE 5% IN PLASTIC CONTAINER, MAGNESIUM SULFATE
 MAGNESIUM SULFATE IN PLASTIC CONTAINER, MAGNESIUM SULFATE
 MEROPENEM, MEROPENEM

HQ SPECLT PHARMA

* HQ SPECIALTY PHARMA CORP
 AMPICILLIN AND SULBACTAM, AMPICILLIN SODIUM
 AMPICILLIN SODIUM, AMPICILLIN SODIUM
 PENICILLIN G POTASSIUM, PENICILLIN G POTASSIUM

HRA PHARMA

* HRA PHARMA RARE DISEASES
 LYSODREN, MITOTANE
 METOPIRONE, METYRAPONE

HUMANWELL

* HUMANWELL PHARMACEUTICAL US INC
 GABAPENTIN, GABAPENTIN

HUMANWELL PURACAP

* HUMANWELL PURACAP PHARMACEUTICAL WUHAN CO LTD
 AMANTADINE HYDROCHLORIDE, AMANTADINE HYDROCHLORIDE
 DUTASTERIDE, DUTASTERIDE
 IBUPROFEN, IBUPROFEN (OTC)
 ICOSAPENT ETHYL, ICOSAPENT ETHYL

HUONS

* HUONS CO LTD
 BUPIVACAINE HYDROCHLORIDE, BUPIVACAINE HYDROCHLORIDE
 LIDOCAINE HYDROCHLORIDE, LIDOCAINE HYDROCHLORIDE

ROCHE

* HOFFMANN LA ROCHE INC
 FUZEON, ENFUVIRTIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** H ****

* HOFFMANN LA ROCHE INC
TAMIFLU, OSELTAMIVIR PHOSPHATE

SHUANGCHENG

* HAINAN SHUANGCHENG PHARMACEUTICALS CO LTD
BIVALIRUDIN, BIVALIRUDIN
EPTIFIBATIDE, EPTIFIBATIDE

**** I ******I 3 PHARMS**

* I 3 PHARMACEUTICALS LLC
GUANFACINE HYDROCHLORIDE, GUANFACINE HYDROCHLORIDE
MARAVIROC, MARAVIROC
POSACONAZOLE, POSACONAZOLE

I3 PHARMS

* I3 PHARMACEUTICALS LLC
ISOSORBIDE DINITRATE AND HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
RAMELTEON, RAMELTEON
RANOLAZINE, RANOLAZINE
SELEGILINE HYDROCHLORIDE, SELEGILINE HYDROCHLORIDE

IBSA

* IBSA INSTITUT BIOCHIMIQUE SA
FLECTOR, DICLOFENAC EPOLAMINE
LIDOCAINE, LIDOCAINE
TIROSINT, LEVOTHYROXINE SODIUM
TIROSINT-SOL, LEVOTHYROXINE SODIUM

IBSA INST BIO

* IBSA INSTITUT BIOCHIMIQUE SA
LICART, DICLOFENAC EPOLAMINE

ICU MEDICAL INC

* ICU MEDICAL INC
ACETIC ACID 0.25% IN PLASTIC CONTAINER, ACETIC ACID, GLACIAL
AMINOSYN II 10% IN PLASTIC CONTAINER, AMINO ACIDS
AMINOSYN II 15% IN PLASTIC CONTAINER, AMINO ACIDS
AMINOSYN-PF 10%, AMINO ACIDS
AMINOSYN-PF 7%, AMINO ACIDS
DEXTROSE 10% IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 20% IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 30% IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 40% IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER, CALCIUM CHLORIDE
DEXTROSE 5% AND SODIUM CHLORIDE 0.225% IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 5% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 50% IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 70% IN PLASTIC CONTAINER, DEXTROSE
GLYCINE 1.5% IN PLASTIC CONTAINER, GLYCINE
IONOSOL MB AND DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
LACTATED RINGER'S IN PLASTIC CONTAINER, CALCIUM CHLORIDE
MANNITOL 20% IN PLASTIC CONTAINER, MANNITOL
NORMOSOL-M AND DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
NORMOSOL-R AND DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
NORMOSOL-R IN PLASTIC CONTAINER, MAGNESIUM CHLORIDE
PHYSIOSOL IN PLASTIC CONTAINER, MAGNESIUM CHLORIDE
POTASSIUM CHLORIDE 0.149% IN SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, POTASSIUM
POTASSIUM CHLORIDE 10MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER,
POTASSIUM CHLORIDE 10MEQ IN PLASTIC CONTAINER, POTASSIUM CHLORIDE
POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER,
POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.225% IN PLASTIC
POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER,
POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,
POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE 20MEQ IN PLASTIC CONTAINER, POTASSIUM CHLORIDE
POTASSIUM CHLORIDE 20MEQ IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, POTASSIUM

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** I ******* ICU MEDICAL INC**

POTASSIUM CHLORIDE 30MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 40MEQ IN PLASTIC CONTAINER, POTASSIUM CHLORIDE
 POTASSIUM CHLORIDE 40MEQ IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, POTASSIUM
 RINGER'S IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, SODIUM CHLORIDE
 SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, SODIUM CHLORIDE
 SORBITOL-MANNITOL IN PLASTIC CONTAINER, MANNITOL
 STERILE WATER FOR INJECTION IN PLASTIC CONTAINER, STERILE WATER FOR INJECTION
 STERILE WATER IN PLASTIC CONTAINER, STERILE WATER FOR IRRIGATION

IDENTIRX*** IDENTIRX LLC**

CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE

IDORSIA*** IDORSIA PHARMACEUTICALS LTD**

QUVIVIQ, DARIDOREXANT HYDROCHLORIDE
 TRYVIO, APROCITENTAN

IMPAX*** IMPAX LABORATORIES LLC**

ADRENACLICK, EPINEPHRINE
 CREXONT, CARBIDOPA
 RYTARY, CARBIDOPA
 SEVELAMER CARBONATE, SEVELAMER CARBONATE

IMPAX LABS*** IMPAX LABORATORIES INC**

ACARBOSE, ACARBOSE
 ANAGRELIDE HYDROCHLORIDE, ANAGRELIDE HYDROCHLORIDE
 BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
 CARBIDOPA AND LEVODOPA, CARBIDOPA
 CHLOROQUINE PHOSPHATE, CHLOROQUINE PHOSPHATE
 CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
 COLESTIPOL HYDROCHLORIDE, COLESTIPOL HYDROCHLORIDE
 DANTROLENE SODIUM, DANTROLENE SODIUM
 DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE
 DIPYRIDAMOLE, DIPYRIDAMOLE
 FENOFIBRATE, FENOFIBRATE
 FLUDROCORTISONE ACETATE, FLUDROCORTISONE ACETATE
 METHYLTESTOSTERONE, METHYLTESTOSTERONE
 MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE
 OMEPRAZOLE, OMEPRAZOLE
 OXYMORPHONE HYDROCHLORIDE, OXYMORPHONE HYDROCHLORIDE
 PILOCARPINE HYDROCHLORIDE, PILOCARPINE HYDROCHLORIDE
 PYRIDOSTIGMINE BROMIDE, PYRIDOSTIGMINE BROMIDE
 RILUZOLE, RILUZOLE
 RIMANTADINE HYDROCHLORIDE, RIMANTADINE HYDROCHLORIDE
 TAMSULOSIN HYDROCHLORIDE, TAMSULOSIN HYDROCHLORIDE
 TERBUTALINE SULFATE, TERBUTALINE SULFATE

IMPAX LABS INC*** IMPAX LABORATORIES INC**

ACITRETIN, ACITRETIN
 BUDESONIDE, BUDESONIDE
 BUSPIRONE HYDROCHLORIDE, BUSPIRONE HYDROCHLORIDE
 CARVEDILOL PHOSPHATE, CARVEDILOL PHOSPHATE
 COLESEVELAM HYDROCHLORIDE, COLESEVELAM HYDROCHLORIDE
 DEXEDRINE SPANSULE, DEXTROAMPHETAMINE SULFATE
 DEXMETHYLPHENIDATE HYDROCHLORIDE, DEXMETHYLPHENIDATE HYDROCHLORIDE
 EMVERM, MEBENDAZOLE
 EPIRUBICIN HYDROCHLORIDE, EPIRUBICIN HYDROCHLORIDE
 FENOFIBRIC ACID, CHOLINE FENOFIBRATE
 GLYBURIDE AND METFORMIN HYDROCHLORIDE, GLYBURIDE
 GLYBURIDE, GLYBURIDE
 HYDROCORTISONE, HYDROCORTISONE
 HYDROXYZINE PAMOATE, HYDROXYZINE PAMOATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** I ******* IMPAX LABORATORIES INC**

LAMOTRIGINE, LAMOTRIGINE
 LEVALBUTEROL HYDROCHLORIDE, LEVALBUTEROL HYDROCHLORIDE
 METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
 METHYLTESTOSTERONE, METHYLTESTOSTERONE
 METOCLOPRAMIDE HYDROCHLORIDE, METOCLOPRAMIDE HYDROCHLORIDE
 NITROFURANTOIN, NITROFURANTOIN, MACROCRYSTALLINE
 PROPRANOLOL HYDROCHLORIDE, PROPRANOLOL HYDROCHLORIDE
 PYRIDOSTIGMINE BROMIDE, PYRIDOSTIGMINE BROMIDE

IMPAX PHARMS

* IMPAX PHARMACEUTICALS
 GEMFIBROZIL, GEMFIBROZIL
 MIDODRINE HYDROCHLORIDE, MIDODRINE HYDROCHLORIDE

IMPEL PHARMS

* IMPEL PHARMACEUTICALS LLC
 TRUDHESA, DIHYDROERGOTAMINE MESYLATE

IMS LTD

* INTERNATIONAL MEDICATION SYSTEMS LTD
 REGADENOSON, REGADENOSON

INCYTE CORP

* INCYTE CORP
 JAKAFI, RUXOLITINIB PHOSPHATE
 OPZELURA, RUXOLITINIB PHOSPHATE
 PEMAZYRE, PEMIGATINIB

INDCHEMIE HEALTH

* INDCHEMIE HEALTH SPECIALTIES PVT LTD
 ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM
 NEBIVOLOL HYDROCHLORIDE, NEBIVOLOL HYDROCHLORIDE

INDICUS PHARMA

* INDICUS PHARMA LLC
 ACETAZOLAMIDE, ACETAZOLAMIDE
 DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
 GLYCOPYRROLATE, GLYCOPYRROLATE

INDIES PHARMA

* INDIES PHARMA JAMAICA LTD
 REGADENOSON, REGADENOSON

INDIVIOR

* INDIVIOR INC
 OPVEE, NALMEFENE HYDROCHLORIDE
 PERSERIS KIT, RISPERIDONE
 SUBLOCADE, BUPRENORPHINE
 SUBOXONE, BUPRENORPHINE HYDROCHLORIDE

INDOCO

* INDOCO REMEDIES LTD
 ACETYLCYSTEINE, ACETYLCYSTEINE
 ALLOPURINOL, ALLOPURINOL
 APIXABAN, APIXABAN
 BRIMONIDINE TARTRATE, BRIMONIDINE TARTRATE
 CETIRIZINE HYDROCHLORIDE, CETIRIZINE HYDROCHLORIDE (OTC)
 DORZOLAMIDE HYDROCHLORIDE AND TIMOLOL MALEATE, DORZOLAMIDE HYDROCHLORIDE
 DORZOLAMIDE HYDROCHLORIDE, DORZOLAMIDE HYDROCHLORIDE
 FEBUXOSTAT, FEBUXOSTAT
 LACOSAMIDE, LACOSAMIDE
 LOFEXIDINE HYDROCHLORIDE, LOFEXIDINE HYDROCHLORIDE
 RASAGILINE MESYLATE, RASAGILINE MESYLATE
 VARENICLINE TARTRATE, VARENICLINE TARTRATE

INDOCO REMEDIES

* INDOCO REMEDIES LTD
 GLIMEPIRIDE, GLIMEPIRIDE

INFORLIFE

* INFORLIFE SA
 ACETAMINOPHEN, ACETAMINOPHEN
 CIPROFLOXACIN IN DEXTROSE 5% IN PLASTIC CONTAINER, CIPROFLOXACIN
 FLUCONAZOLE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, FLUCONAZOLE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** I ******* INFORLIFE SA**

LEVOFLOXACIN IN DEXTROSE 5% IN PLASTIC CONTAINER, LEVOFLOXACIN
 METRONIDAZOLE IN PLASTIC CONTAINER, METRONIDAZOLE
 MIDAZOLAM IN 0.9% SODIUM CHLORIDE, MIDAZOLAM
 NICARDIPINE HYDROCHLORIDE IN 0.83% SODIUM CHLORIDE, NICARDIPINE HYDROCHLORIDE
 NICARDIPINE HYDROCHLORIDE IN 0.86% SODIUM CHLORIDE, NICARDIPINE HYDROCHLORIDE
 NOREPINEPHRINE BITARTRATE IN 0.9% SODIUM CHLORIDE, NOREPINEPHRINE BITARTRATE
 ROPIVACAINE HYDROCHLORIDE, ROPIVACAINE HYDROCHLORIDE
 ZOLEDRONIC ACID, ZOLEDRONIC ACID

INGENUS PHARMS LLC*** INGENUS PHARMACEUTICALS LLC**

ARSENIC TRIOXIDE, ARSENIC TRIOXIDE
 CABERGOLINE, CABERGOLINE
 DORZOLAMIDE HYDROCHLORIDE AND TIMOLOL MALEATE, DORZOLAMIDE HYDROCHLORIDE
 IVABRADINE HYDROCHLORIDE, IVABRADINE HYDROCHLORIDE
 LEVETIRACETAM, LEVETIRACETAM
 MELPHALAN HYDROCHLORIDE, MELPHALAN HYDROCHLORIDE
 PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
 TAFLUPROST, TAFLUPROST
 TIMOLOL MALEATE, TIMOLOL MALEATE

INNOGENIX*** INNOGENIX LLC**

HALOPERIDOL, HALOPERIDOL
 METOLAZONE, METOLAZONE
 METRONIDAZOLE, METRONIDAZOLE
 PILOCARPINE HYDROCHLORIDE, PILOCARPINE HYDROCHLORIDE
 PROPRANOLOL HYDROCHLORIDE, PROPRANOLOL HYDROCHLORIDE

INNOPHARMA*** INNOPHARMA LICENSING LLC A SUB OF PFIZER INC**

PROPOFOL, PROPOFOL

INSMED INC*** INSMED INC**

ARIKAYCE KIT, AMIKACIN SULFATE

INTAS PHARMS USA*** INTAS PHARMACEUTICALS USA**

IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE

INTELLIPHARMACEUTICS*** INTELLIPHARMACEUTICS CORP**

DESVENLAFAXINE SUCCINATE, DESVENLAFAXINE SUCCINATE
 DEXMETHYLPHENIDATE HYDROCHLORIDE, DEXMETHYLPHENIDATE HYDROCHLORIDE
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 QUETIAPINE FUMARATE, QUETIAPINE FUMARATE
 VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE

INTERCEPT PHARMS INC*** INTERCEPT PHARMACEUTICALS INC**

OICALIVA, OBETICHOLIC ACID

INTERSECT ENT INC*** INTERSECT ENT INC**

SINUVA, MOMETASONE FUROATE

INTL ISOTOPES*** INTERNATIONAL ISOTOPES INC**

SODIUM IODIDE I 131, SODIUM IODIDE I-131

INTL MEDICATION*** INTERNATIONAL MEDICATION SYSTEM**

LARYNG-O-JET KIT, LIDOCAINE HYDROCHLORIDE
 LIDOCAINE HYDROCHLORIDE, LIDOCAINE HYDROCHLORIDE
 NALOXONE HYDROCHLORIDE, NALOXONE HYDROCHLORIDE
 PHYTONADIONE, PHYTONADIONE
 PROCAINAMIDE HYDROCHLORIDE, PROCAINAMIDE HYDROCHLORIDE

INTL MEDICATION SYS*** INTERNATIONAL MEDICATION SYSTEMS LTD**

ATROPINE SULFATE, ATROPINE SULFATE
 CALCIUM CHLORIDE 10%, CALCIUM CHLORIDE
 DEXTROSE 50%, DEXTROSE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** I ****

* INTERNATIONAL MEDICATION SYSTEMS LTD
 EPINEPHRINE, EPINEPHRINE
 LORAZEPAM, LORAZEPAM
 MORPHINE SULFATE, MORPHINE SULFATE
 SODIUM BICARBONATE, SODIUM BICARBONATE

INTRA-CELLULAR

* INTRA-CELLULAR THERAPIES INC
 CAPLYTA, LUMATEPERONE TOSYLATE

INTRABIO

* INTRABIO INC
 AQNEURSA, LEVACETYLLEUCINE

INVAGEN PHARMS

* INVAGEN PHARMACEUTICALS INC
 ALFUZOSIN HYDROCHLORIDE, ALFUZOSIN HYDROCHLORIDE
 AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
 CALCIUM ACETATE, CALCIUM ACETATE
 CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE
 CYCLOBENZAPRINE HYDROCHLORIDE, CYCLOBENZAPRINE HYDROCHLORIDE
 ESCITALOPRAM OXALATE, ESCITALOPRAM OXALATE
 FOSINOPRIL SODIUM AND HYDROCHLOROTHIAZIDE, FOSINOPRIL SODIUM
 FOSINOPRIL SODIUM, FOSINOPRIL SODIUM
 GABAPENTIN, GABAPENTIN
 GEMFIBROZIL, GEMFIBROZIL
 LANREOTIDE ACETATE, LANREOTIDE ACETATE
 LANTHANUM CARBONATE, LANTHANUM CARBONATE
 LEUPROLIDE ACETATE FOR DEPOT SUSPENSION, LEUPROLIDE ACETATE
 LINAGLIPTIN, LINAGLIPTIN
 LISINOPRIL AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 LISINOPRIL, LISINOPRIL
 LURASIDONE HYDROCHLORIDE, LURASIDONE HYDROCHLORIDE
 MEPROBAMATE, MEPROBAMATE
 NABUMETONE, NABUMETONE
 NADOLOL, NADOLOL
 OSELTAMIVIR PHOSPHATE, OSELTAMIVIR PHOSPHATE
 PENICILLAMINE, PENICILLAMINE
 PREGABALIN, PREGABALIN
 RALOXIFENE HYDROCHLORIDE, RALOXIFENE HYDROCHLORIDE
 SERTRALINE HYDROCHLORIDE, SERTRALINE HYDROCHLORIDE
 SEVELAMER CARBONATE, SEVELAMER CARBONATE
 SILDENAFIL CITRATE, SILDENAFIL CITRATE
 SIMVASTATIN, SIMVASTATIN
 TERBINAFINE HYDROCHLORIDE, TERBINAFINE HYDROCHLORIDE
 TOPIRAMATE, TOPIRAMATE
 TROSPIUM CHLORIDE, TROSPIUM CHLORIDE
 VIGABATRIN, VIGABATRIN
 VILAZODONE HYDROCHLORIDE, VILAZODONE HYDROCHLORIDE
 WARFARIN SODIUM, WARFARIN SODIUM
 ZONISAMIDE, ZONISAMIDE

INVATECH

* INVATECH PHARMA SOLUTIONS LLC
 DIVALPROEX SODIUM, DIVALPROEX SODIUM
 MECLIZINE HYDROCHLORIDE, MECLIZINE HYDROCHLORIDE

INVENTIA

* INVENTIA HEALTHCARE LTD
 CHLORTHALIDONE, CHLORTHALIDONE
 COLESEVELAM HYDROCHLORIDE, COLESEVELAM HYDROCHLORIDE
 DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE
 LANSOPRAZOLE, LANSOPRAZOLE
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 NITROFURANTOIN (MONOHYDRATE/MACROCRYSTALS), NITROFURANTOIN
 OLMESARTAN MEDOXOMIL, OLMESARTAN MEDOXOMIL
 PALIPERIDONE, PALIPERIDONE
 TELMISARTAN, TELMISARTAN

INVENTIA HLTHCARE

* INVENTIA HEALTHCARE PRIVATE LTD

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** I ****

* INVENTIA HEALTHCARE PRIVATE LTD

BUSPIRONE HYDROCHLORIDE, BUSPIRONE HYDROCHLORIDE
 FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 TOLTERODINE TARTRATE, TOLTERODINE TARTRATE
 VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE

IONETIX

* IONETIX CORP

AMMONIA N 13, AMMONIA N-13

IONIS PHARMS INC

* IONIS PHARMACEUTICALS INC

TRYNGOLZA (AUTOINJECTOR), OLEZARSEN SODIUM

IPCA LABS

* IPCA LABORATORIES LTD

LAMOTRIGINE, LAMOTRIGINE

IPCA LABS LTD

* IPCA LABORATORIES LTD

ALENDRONATE SODIUM, ALENDRONATE SODIUM
 CHLORTHALIDONE, CHLORTHALIDONE
 ETODOLAC, ETODOLAC
 FUROSEMIDE, FUROSEMIDE
 HYDROXYCHLOROQUINE SULFATE, HYDROXYCHLOROQUINE SULFATE
 METOCLOPRAMIDE HYDROCHLORIDE, METOCLOPRAMIDE HYDROCHLORIDE
 METOPROLOL TARTRATE, METOPROLOL TARTRATE
 PROPRANOLOL HYDROCHLORIDE, PROPRANOLOL HYDROCHLORIDE
 RISPERIDONE, RISPERIDONE

IPR

* IPR PHARMACEUTICALS INC

CRESTOR, ROSUVASTATIN CALCIUM

IPSEN

* IPSEN BIOPHARMACEUTICALS INC

BYLVAY, ODEVIXIBAT
 IQIRVO, ELAFIBRANOR
 ONIVYDE, IRINOTECAN HYDROCHLORIDE
 SOHONOS, PALOVAROTENE

IPSEN PHARMA

* IPSEN PHARMA BIOTECH SAS

SOMATULINE DEPOT, LANREOTIDE ACETATE

IRONSHORE PHARMS

* IRONSHORE PHARMACEUTICALS INC A WHOLLY OWNED SUB OF COLLEGIUM PHARMACEUTICAL INC

JORNAY PM, METHYLPHENIDATE HYDROCHLORIDE

ISOLOGIC INNOVATIVE

* ISOLOGIC INNOVATIVE RADIOPHARMACEUTICALS LTD

FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18

ISOTEX

* ISOTEX DIAGNOSTICS

GLOFIL-125, IOTHALAMATE SODIUM I-125

ISTITUTO BIO ITA SPA

* ISTITUTO BIOCHIMICO ITALIANO SPA

AMPICILLIN AND SULBACTAM, AMPICILLIN SODIUM
 AMPICILLIN SODIUM, AMPICILLIN SODIUM
 NAFCILLIN SODIUM, NAFCILLIN SODIUM
 PENICILLIN G POTASSIUM, PENICILLIN G POTASSIUM
 PIPERACILLIN AND TAZOBACTAM, PIPERACILLIN SODIUM
 PIPERACILLIN, PIPERACILLIN SODIUM

ITALFARMACO SA

* ITALFARMACO SA

TIGLUTIK KIT, RILUZOLE

ITALFARMACO SPA

* ITALFARMACO SPA

DUVYZAT, GIVINOSTAT HYDROCHLORIDE

ITERUM THERAP

* ITERUM THERAPEUTICS US LTD

ORLYNVAH, PROBENECID

IVAX SUB TEVA PHARMS

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** I ****

* IVAX PHARMACEUTICALS INC SUB TEVA PHARMACEUTICALS USA
 ANAGRELIDE HYDROCHLORIDE, ANAGRELIDE HYDROCHLORIDE
 BACLOFEN, BACLOFEN
 CABERGOLINE, CABERGOLINE
 CETIRIZINE HYDROCHLORIDE AND PSEUDOEPHEDRINE HYDROCHLORIDE, CETIRIZINE HYDROCHLORIDE
 CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE
 CLOZAPINE, CLOZAPINE
 CYCLOSPORINE, CYCLOSPORINE
 DIAZEPAM, DIAZEPAM
 ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM
 FAMOTIDINE, FAMOTIDINE
 FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 GABAPENTIN, GABAPENTIN
 HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 TOLTERODINE TARTRATE, TOLTERODINE TARTRATE

**** J ******HENGRUI PHARMA**

* JIANGSU HENGRUI PHARMACEUTICALS CO LTD
 BUPIVACAINE LIPOSOME, BUPIVACAINE
 CARMUSTINE, CARMUSTINE
 CASPOFUNGIN ACETATE, CASPOFUNGIN ACETATE
 CISATRACURIUM BESYLATE PRESERVATIVE FREE, CISATRACURIUM BESYLATE
 CISATRACURIUM BESYLATE, CISATRACURIUM BESYLATE
 CYCLOPHOSPHAMIDE, CYCLOPHOSPHAMIDE
 DAPTOMYCIN, DAPTOMYCIN
 DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE
 DOCETAXEL, DOCETAXEL
 FONDAPARINUX SODIUM, FONDAPARINUX SODIUM
 GADOBUTROL, GADOBUTROL
 GADOTERATE MEGLUMINE, GADOTERATE MEGLUMINE
 IODIXANOL, IODIXANOL
 IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE
 OXALIPLATIN, OXALIPLATIN
 PACLITAXEL, PACLITAXEL
 THIOTEPA, THIOTEPA

J AND J CONSUMER INC

* JOHNSON AND JOHNSON CONSUMER INC MCNEIL CONSUMER HEALTHCARE DIV
 PEPCID AC, FAMOTIDINE (OTC)

JANSSEN BIOTECH

* JANSSEN BIOTECH INC
 AKEEGA, ABIRATERONE ACETATE
 BALVERSA, ERDAFITINIB
 ERLEADA, APALUTAMIDE
 LAZCLUZE, LAZERTINIB MESYLATE
 ZYTIGA, ABIRATERONE ACETATE

JANSSEN PHARMS

* JANSSEN PHARMACEUTICALS INC
 CONCERTA, METHYLPHENIDATE HYDROCHLORIDE
 ELMIRON, PENTOSAN POLYSULFATE SODIUM
 HALDOL, HALOPERIDOL DECANOATE
 INVEGA HAFYERA, PALIPERIDONE PALMITATE
 INVEGA SUSTENNA, PALIPERIDONE PALMITATE
 INVEGA TRINZA, PALIPERIDONE PALMITATE
 INVEGA, PALIPERIDONE
 INVOKAMET XR, CANAGLIFLOZIN
 INVOKAMET, CANAGLIFLOZIN
 INVOKANA, CANAGLIFLOZIN
 RISPERDAL CONSTA, RISPERIDONE
 RISPERDAL, RISPERIDONE
 SPORANOX, ITRACONAZOLE
 SPRAVATO, ESKETAMINE HYDROCHLORIDE
 TOPAMAX, TOPIRAMATE
 XARELTO, RIVAROXABAN

JANSSEN PRODS

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** J ******* JANSSEN PRODUCTS LP**

EDURANT, RILPIVIRINE HYDROCHLORIDE
 PREZCOBIX, COBICISTAT
 PREZISTA, DARUNAVIR
 SYMTUZA, COBICISTAT
 YONDELIS, TRABECTEDIN

JANSSEN R AND D

* JANSSEN RESEARCH AND DEVELOPMENT LLC
 INTELENCE, ETRAVIRINE

JANSSEN THERAP

* JANSSEN THERAPEUTICS DIV JANSSEN PRODUCTS LP
 SIRTURO, BEDAQUILINE FUMARATE

JAZZ

* JAZZ PHARMACEUTICALS IRELAND LTD
 XYWAV, CALCIUM OXYBATE
 ZEPZELCA, LURBINECTEDIN

JAZZ PHARMS

* JAZZ PHARMACEUTICALS INC
 XYREM, SODIUM OXYBATE

JAZZ PHARMS INC

* JAZZ PHARMACEUTICALS INC
 DEFITELIO, DEFIBROTIDE SODIUM

JAZZ PHARMS RES

* JAZZ PHARMACEUTICALS RESEARCH UK LTD
 EPIDIOLEX, CANNABIDIOL

JDP

* JDP THERAPEUTICS LLC
 QUZYTIR, CETIRIZINE HYDROCHLORIDE

JIANGSU HANSOH PHARM

* JIANGSU HANSOH PHARMACEUTICAL GROUP CO LTD
 AZACITIDINE, AZACITIDINE
 BORTEZOMIB, BORTEZOMIB
 DECITABINE, DECITABINE
 FULVESTRANT, FULVESTRANT
 GEMCITABINE HYDROCHLORIDE, GEMCITABINE HYDROCHLORIDE
 ICATIBANT ACETATE, ICATIBANT ACETATE
 MICAFUNGIN SODIUM, MICAFUNGIN SODIUM
 PEMETREXED DISODIUM, PEMETREXED DISODIUM
 VINOELBINE TARTRATE, VINOELBINE TARTRATE

JIANGXI QINGFENG

* JIANGXI QINGFENG PHARMACEUTICAL CO LTD
 ERIBULIN MESYLATE, ERIBULIN MESYLATE

JOHNS HOPKINS UNIV

* JOHNS HOPKINS UNIV
 AMMONIA N 13, AMMONIA N-13

JOURNEY

* JOURNEY MEDICAL CORP
 AMZEEQ, MINOCYCLINE HYDROCHLORIDE
 EMROSI, MINOCYCLINE HYDROCHLORIDE
 EURAX, CROTAMITON
 EXELDERM, SULCONAZOLE NITRATE
 QBREXZA, GLYCOPYRRONIUM TOSYLATE
 ZILXI, MINOCYCLINE HYDROCHLORIDE

JUBILANT

* JUBILANT DRAXIMAGE INC DBA JUBILANT RADIOPHARMA
 DRAX EXAMETAZIME, TECHNETIUM TC-99M EXAMETAZIME KIT
 DRAXIMAGE DTPA, TECHNETIUM TC-99M PENTETATE KIT
 DRAXIMAGE MDP-25, TECHNETIUM TC-99M MEDRONATE
 HICON, SODIUM IODIDE I-131
 RUBY-FILL, RUBIDIUM CHLORIDE RB-82
 SODIUM IODIDE I 131, SODIUM IODIDE I-131
 TECHNETIUM TC-99M SULFUR COLLOID KIT, TECHNETIUM TC-99M SULFUR COLLOID KIT

JUBILANT CADISTA

* JUBILANT CADISTA PHARMACEUTICALS INC

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** J ****

* JUBILANT CADISTA PHARMACEUTICALS INC
 CLOMIPRAMINE HYDROCHLORIDE, CLOMIPRAMINE HYDROCHLORIDE
 CYCLOBENZAPRINE HYDROCHLORIDE, CYCLOBENZAPRINE HYDROCHLORIDE
 DOXEPIN HYDROCHLORIDE, DOXEPIN HYDROCHLORIDE
 ESCITALOPRAM OXALATE, ESCITALOPRAM OXALATE
 HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 LAMOTRIGINE, LAMOTRIGINE
 LOPERAMIDE HYDROCHLORIDE, LOPERAMIDE HYDROCHLORIDE
 LOSARTAN POTASSIUM AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 LOSARTAN POTASSIUM, LOSARTAN POTASSIUM
 MECLIZINE HYDROCHLORIDE, MECLIZINE HYDROCHLORIDE
 METHENAMINE HIPPURATE, METHENAMINE HIPPURATE
 METHYLPREDNISOLONE, METHYLPREDNISOLONE
 PREDNISONE, PREDNISONE
 PROCOMP, PROCHLORPERAZINE MALEATE
 TERAZOSIN HYDROCHLORIDE, TERAZOSIN HYDROCHLORIDE

JUBILANT DRAXIMAGE

* JUBILANT DRAXIMAGE INC
 TECHNETIUM TC 99M MERTIATIDE KIT, TECHNETIUM TC-99M MERTIATIDE KIT

* JUBILANT DRAXIMAGE RADIOPHARMACIES INC
 FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18
 SODIUM FLUORIDE F-18, SODIUM FLUORIDE F-18

* JUBILANT DRAXIMAGE USA INC
 TECHNETIUM TC 99M SESTAMIBI, TECHNETIUM TC-99M SESTAMIBI KIT

JUBILANT GENERICS

* JUBILANT GENERICS LTD
 CHILDREN'S CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 CHILDREN'S CETIRIZINE HYDROCHLORIDE HIVES RELIEF, CETIRIZINE HYDROCHLORIDE (OTC)
 CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
 DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
 IRBESARTAN, IRBESARTAN
 LURASIDONE HYDROCHLORIDE, LURASIDONE HYDROCHLORIDE
 OLANZAPINE, OLANZAPINE
 RISPERIDONE, RISPERIDONE
 SILDENAFIL CITRATE, SILDENAFIL CITRATE
 SOLIFENACIN SUCCINATE, SOLIFENACIN SUCCINATE
 SPIRONOLACTONE, SPIRONOLACTONE
 TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE
 VALACYCLOVIR HYDROCHLORIDE, VALACYCLOVIR HYDROCHLORIDE
 VALSARTAN, VALSARTAN
 ZOLMITRIPTAN, ZOLMITRIPTAN

STEVENS J

* JEROME STEVENS PHARMACEUTICALS INC
 BUTALBITAL, ASPIRIN, CAFFEINE, AND CODEINE PHOSPHATE, ASPIRIN
 DIGOXIN, DIGOXIN
 UNITHROID, LEVOTHYROXINE SODIUM **

**** K ******GRIFFEN**

* KW GRIFFEN CO
 BIOSCRUB, CHLORHEXIDINE GLUCONATE (OTC)

KADMON PHARMS LLC

* KADMON PHARMACEUTICALS LLC
 REZUROCK, BELUMOSUDIL MESYLATE

KAI PHARMS INC

* KAI PHARMACEUTICALS INC A WHOLLY OWNED SUBSIDIARY OF AMGEN INC
 PARSABIV, ETELCALCETIDE

KALEO INC

* KALEO INC
 AUVI-Q, EPINEPHRINE

KAMAT

* KAMAT PHARMATECH LLC
 RALDESY, TRAZODONE HYDROCHLORIDE

KANCHAN HLTHCARE

* KANCHAN HEALTHCARE INC

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** K ****

* KANCHAN HEALTHCARE INC
 CYCLOPHOSPHAMIDE, CYCLOPHOSPHAMIDE
 DEXAMETHASONE, DEXAMETHASONE
 LACOSAMIDE, LACOSAMIDE
 MYAMBUTOL, ETHAMBUTOL HYDROCHLORIDE
 OXCARBAZEPINE, OXCARBAZEPINE

KARO HLTHCARE

* KARO HEALTHCARE INC
 LAMISIL AT, TERBINAFINE (OTC)
 LAMISIL AT, TERBINAFINE HYDROCHLORIDE (OTC)
 LAMISIL, TERBINAFINE HYDROCHLORIDE (OTC)

KARYOPHARM THERAPS

* KARYOPHARM THERAPEUTICS INC
 XPOVIO, SELINEXOR

KENTON

* KENTON CHEMICALS AND PHARMACEUTICALS CORP
 ANASTROZOLE, ANASTROZOLE
 CYPROHEPTADINE HYDROCHLORIDE, CYPROHEPTADINE HYDROCHLORIDE
 RILUZOLE, RILUZOLE

KENVUE BRANDS

* KENVUE BRANDS LLC
 CHILDREN'S MOTRIN COLD, IBUPROFEN (OTC)
 CHILDREN'S MOTRIN, IBUPROFEN (OTC)
 CHILDREN'S ZYRTEC ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 CHILDREN'S ZYRTEC HIVES, CETIRIZINE HYDROCHLORIDE (OTC)
 IMODIUM A-D, LOPERAMIDE HYDROCHLORIDE (OTC)
 IMODIUM MULTI-SYMPTOM RELIEF, LOPERAMIDE HYDROCHLORIDE (OTC)
 JUNIOR STRENGTH MOTRIN, IBUPROFEN (OTC)
 MEN'S ROGAINE, MINOXIDIL (OTC)
 MOTRIN IB, IBUPROFEN (OTC)
 PEPCID AC, FAMOTIDINE (OTC)
 PEPCID COMPLETE, CALCIUM CARBONATE (OTC)
 RHINOCORT ALLERGY, BUDESONIDE (OTC)
 ROGAINE (FOR MEN), MINOXIDIL (OTC)
 ROGAINE (FOR WOMEN), MINOXIDIL (OTC)
 ROGAINE EXTRA STRENGTH (FOR MEN), MINOXIDIL (OTC)
 SINE-AID IB, IBUPROFEN (OTC)
 SUDAFED 24 HOUR, PSEUDOEPHEDRINE HYDROCHLORIDE (OTC)
 TYLENOL, ACETAMINOPHEN (OTC)
 VISINE L.R., OXYMETAZOLINE HYDROCHLORIDE (OTC)
 VISINE, NAPHAZOLINE HYDROCHLORIDE (OTC)
 WOMEN'S ROGAINE, MINOXIDIL (OTC)
 ZYRTEC ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 ZYRTEC-D 12 HOUR, CETIRIZINE HYDROCHLORIDE (OTC)

KERYX BIOPHARMS

* KERYX BIOPHARMACEUTICALS INC
 AURYXIA, FERRIC CITRATE

KETTERING MEDCTR

* KETTERING MEDCTR
 FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18

KEY THERAP

* KEY THERAPEUTICS LLC
 CETRAXAL, CIPROFLOXACIN HYDROCHLORIDE

KINDEVA

* KINDEVA DRUG DELIVERY LP
 FENTANYL-100, FENTANYL
 FENTANYL-12, FENTANYL
 FENTANYL-25, FENTANYL
 FENTANYL-50, FENTANYL
 FENTANYL-75, FENTANYL
 PROVENTIL-HFA, ALBUTEROL SULFATE

KINDOS

* KINDOS PHARMACEUTICALS CO LTD
 DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
 FLUOROURACIL, FLUOROURACIL

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** K ****

* KINDOS PHARMACEUTICALS CO LTD
ROPIVACAINE HYDROCHLORIDE, ROPIVACAINE HYDROCHLORIDE

KING PHARMS

* KING PHARMACEUTICALS RESEARCH AND DEVELOPMENT LLC
CYTOMEL, LIOTHYRONINE SODIUM
LEVOXYL, LEVOTHYROXINE SODIUM **

KING PHARMS LLC

* KING PHARMACEUTICALS LLC
ALTACE, RAMIPRIL
BICILLIN C-R 900/300, PENICILLIN G BENZATHINE
BICILLIN C-R, PENICILLIN G BENZATHINE
BICILLIN L-A, PENICILLIN G BENZATHINE
PENICILLIN G PROCAINE, PENICILLIN G PROCAINE
SILVADENE, SILVER SULFADIAZINE

KNACK

* KNACK PHARMACEUTICALS INC
PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM

KNIGHT THERAPS

* KNIGHT THERAPEUTICS USA INC
IMPAVIDO, MILTEFOSINE

KOWA CO

* KOWA CO LTD
LIVALO, PITAVASTATIN CALCIUM

KRAMER

* KRAMER LABORATORIES INC
NIZORAL ANTI-DANDRUFF, KETOCONAZOLE (OTC)

KREITCHMAN PET CTR

* KREITCHMAN PET CENTER
AMMONIA N 13, AMMONIA N-13
FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18
SODIUM FLUORIDE F-18, SODIUM FLUORIDE F-18

KVK TECH

* KVK TECH INC
BENZPHETAMINE HYDROCHLORIDE, BENZPHETAMINE HYDROCHLORIDE
BETAXOLOL HYDROCHLORIDE, BETAXOLOL HYDROCHLORIDE
CYCLOBENZAPRINE HYDROCHLORIDE, CYCLOBENZAPRINE HYDROCHLORIDE
HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE
KALEXATE, SODIUM POLYSTYRENE SULFONATE
PHENDIMETRAZINE TARTRATE, PHENDIMETRAZINE TARTRATE
PHENTERMINE HYDROCHLORIDE, PHENTERMINE HYDROCHLORIDE
PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE

KVK TECH INC

* KVK TECH INC
PHENTERMINE HYDROCHLORIDE, PHENTERMINE HYDROCHLORIDE

KYOWA KIRIN

* KYOWA KIRIN INC
NOURIANZ, ISTRADefylline

**** L ******L PERRIGO CO**

* L PERRIGO CO
ACETAMINOPHEN AND IBUPROFEN, ACETAMINOPHEN (OTC)
CIMETIDINE, CIMETIDINE (OTC)
FEXOFENADINE HYDROCHLORIDE, FEXOFENADINE HYDROCHLORIDE (OTC)
GUAIFENESIN AND PSEUDOEPHEDRINE HYDROCHLORIDE, GUAIFENESIN (OTC)
IBUPROFEN, IBUPROFEN (OTC)
LOPERAMIDE HYDROCHLORIDE, LOPERAMIDE HYDROCHLORIDE (OTC)
MICONAZOLE NITRATE COMBINATION PACK, MICONAZOLE NITRATE (OTC)
MINOXIDIL (FOR MEN), MINOXIDIL (OTC)
MINOXIDIL (FOR WOMEN), MINOXIDIL (OTC)
NICOTINE POLACRILEX, NICOTINE POLACRILEX (OTC)
OMEPRAZOLE MAGNESIUM, OMEPRAZOLE MAGNESIUM (OTC)
PSEUDOEPHEDRINE HYDROCHLORIDE, PSEUDOEPHEDRINE HYDROCHLORIDE (OTC)

LA JOLLA PHARMA

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** L ****

* LA JOLLA PHARMA LLC
GIAPREZA, ANGIOTENSIN II ACETATE

LAB HRA PHARMA

* LABORATOIRE HRA PHARMA
ELLA, ULIPRISTAL ACETATE

LABORATOIRE HRA

* LABORATOIRE HRA PHARMA
LEVONORGESTREL, LEVONORGESTREL (OTC)
OPILL, NORGESTREL (OTC)

LABORATORIOS GRIFOLS

* LABORATORIOS GRIFOLS SA
SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, SODIUM CHLORIDE

LABORATORIOS SALVAT

* LABORATORIOS SALVAT SA
OTOVEL, CIPROFLOXACIN HYDROCHLORIDE

LABS JUVISE

* LABORATOIRES JUVISE PHARMACEUTICALS
PYLERA, BISMUTH SUBCITRATE POTASSIUM

LACER PHARMA

* LACER PHARMA LLC
ERTACZO, SERTACONAZOLE NITRATE

LANDELA PHARM

* LANDELA PHARMACEUTICAL
FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE

LANNETT

* LANNETT CO INC
DICYCLOMINE HYDROCHLORIDE, DICYCLOMINE HYDROCHLORIDE
DIPHENOXYLATE HYDROCHLORIDE AND ATROPINE SULFATE, ATROPINE SULFATE
PRIMIDONE, PRIMIDONE
PROBALAN, PROBENECID

LANNETT CO INC

* LANNETT CO INC
ARIPIPIRAZOLE, ARIPIPIRAZOLE
ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM
BACLOFEN, BACLOFEN
BUPRENORPHINE HYDROCHLORIDE AND NALOXONE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE
BUTALBITAL, ACETAMINOPHEN AND CAFFEINE, ACETAMINOPHEN
CHLORPROMAZINE HYDROCHLORIDE, CHLORPROMAZINE HYDROCHLORIDE
CODEINE SULFATE, CODEINE SULFATE
DANAZOL, DANAZOL
DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE
DIETHYLPROPION HYDROCHLORIDE, DIETHYLPROPION HYDROCHLORIDE
DOXEPIN HYDROCHLORIDE, DOXEPIN HYDROCHLORIDE
DOXYCYCLINE, DOXYCYCLINE
ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM
FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
FLUPHENAZINE HYDROCHLORIDE, FLUPHENAZINE HYDROCHLORIDE
HALOPERIDOL, HALOPERIDOL LACTATE
HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE
LACTULOSE, LACTULOSE
LEVOFLOXACIN, LEVOFLOXACIN
LIDOCAINE HYDROCHLORIDE, LIDOCAINE HYDROCHLORIDE
LISDEXAMFETAMINE DIMESYLATE, LISDEXAMFETAMINE DIMESYLATE
LOPINAVIR AND RITONAVIR, LOPINAVIR
LORATADINE, LORATADINE (OTC)
LOXAPINE SUCCINATE, LOXAPINE SUCCINATE
METAXALONE, METAXALONE
METHADONE HYDROCHLORIDE, METHADONE HYDROCHLORIDE
METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
MONTELUKAST SODIUM, MONTELUKAST SODIUM
MYCOPHENOLATE MOFETIL, MYCOPHENOLATE MOFETIL
NIACIN, NIACIN
OMEPRAZOLE, OMEPRAZOLE
OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE
PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** L ******* LANNETT CO INC**

PAROXETINE HYDROCHLORIDE, PAROXETINE HYDROCHLORIDE
 PILOCARPINE HYDROCHLORIDE, PILOCARPINE HYDROCHLORIDE
 PREDNISOLONE, PREDNISOLONE
 RABEPRAZOLE SODIUM, RABEPRAZOLE SODIUM
 RISPERIDONE, RISPERIDONE
 SUMATRIPTAN, SUMATRIPTAN
 TERBUTALINE SULFATE, TERBUTALINE SULFATE
 TRIAMTERENE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 URSODIOL, URSODIOL

LANTHEUS MEDCL*** LANTHEUS MEDICAL IMAGING INC**

CARDIOLITE, TECHNETIUM TC-99M SESTAMIBI KIT
 DEFINITY RT, PERFLUTREN
 DEFINITY, PERFLUTREN
 NEUROLITE, TECHNETIUM TC-99M BICISATE KIT
 TECHNELITE, TECHNETIUM TC-99M SODIUM PERTECHNETATE GENERATOR
 XENON XE 133, XENON XE-133

LARKEN LABS*** LARKEN LABORATORIES INC**

DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE

LARKEN LABS INC*** LARKEN LABORATORIES INC**

ALLZITAL, ACETAMINOPHEN
 BUTALBITAL AND ACETAMINOPHEN, ACETAMINOPHEN
 DEXAMETHASONE, DEXAMETHASONE

LATINA PHARMA*** LATINA PHARMA SPA**

GLEOSTINE, LOMUSTINE

LAURUS*** LAURUS LABS LTD**

ABACAVIR SULFATE AND LAMIVUDINE, ABACAVIR SULFATE
 ATAZANAVIR SULFATE, ATAZANAVIR SULFATE
 ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM
 EFAVIRENZ, EMTRICITABINE, AND TENOFOVIR DISOPROXIL FUMARATE, EFAVIRENZ
 EFAVIRENZ, LAMIVUDINE AND TENOFOVIR DISOPROXIL FUMARATE, EFAVIRENZ
 EMTRICITABINE AND TENOFOVIR DISOPROXIL FUMARATE, EMTRICITABINE
 GABAPENTIN, GABAPENTIN
 HYDROXYCHLOROQUINE SULFATE, HYDROXYCHLOROQUINE SULFATE
 LOPINAVIR AND RITONAVIR, LOPINAVIR
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 OSELTAMIVIR PHOSPHATE, OSELTAMIVIR PHOSPHATE
 PIRFENIDONE, PIRFENIDONE
 SACUBITRIL AND VALSARTAN, SACUBITRIL

LAVIPHARM*** LAVIPHARM SA**

CATAPRES-TTS-1, CLONIDINE
 CATAPRES-TTS-2, CLONIDINE
 CATAPRES-TTS-3, CLONIDINE

LEADIANT BIOSCI INC*** LEADIANT BIOSCIENCES INC**

ABELCET, AMPHOTERICIN B
 CARNITOR SF, LEVOCARNITINE
 CARNITOR, LEVOCARNITINE
 CYSTARAN, CYSTEAMINE HYDROCHLORIDE
 MATULANE, PROCARBAZINE HYDROCHLORIDE

LEADING*** LEADING PHARMA LLC**

BENZTROPINE MESYLATE, BENZTROPINE MESYLATE
 CARBINOXAMINE MALEATE, CARBINOXAMINE MALEATE
 DIPHENOXYLATE HYDROCHLORIDE AND ATROPINE SULFATE, ATROPINE SULFATE
 DOXEPIN HYDROCHLORIDE, DOXEPIN HYDROCHLORIDE
 FOLIC ACID, FOLIC ACID
 FUROSEMIDE, FUROSEMIDE
 HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** L ****

* LEADING PHARMA LLC
 HYDROXYUREA, HYDROXYUREA
 IMIPRAMINE HYDROCHLORIDE, IMIPRAMINE HYDROCHLORIDE
 KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
 LEUCOVORIN CALCIUM, LEUCOVORIN CALCIUM
 LORAZEPAM, LORAZEPAM
 METYROSINE, METYROSINE
 MEXILETINE HYDROCHLORIDE, MEXILETINE HYDROCHLORIDE
 OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE
 PROCHLORPERAZINE MALEATE, PROCHLORPERAZINE MALEATE

LEGACY PHARMA

* LEGACY PHARMA INC
 ALA-SCALP, HYDROCORTISONE
 BETAPACE AF, SOTALOL HYDROCHLORIDE
 BETAPACE, SOTALOL HYDROCHLORIDE
 BRISDELLE, PAROXETINE MESYLATE
 CLODERM, CLOCORTOLONE PIVALATE
 CROTAN, CROTAMITON
 IMURAN, AZATHIOPRINE
 LOTRONEX, ALOSETRON HYDROCHLORIDE
 MICORT-HC, HYDROCORTISONE ACETATE
 MOTOFEN, ATROPINE SULFATE
 NAFTIN, NAFTIFINE HYDROCHLORIDE
 PRAMOSONE, HYDROCORTISONE ACETATE
 RIDAURA, AURANOFIN

LEGACY PHARMA USA

* LEGACY PHARMA USA INC
 ZANAFLEX, TIZANIDINE HYDROCHLORIDE

LEO PHARMA AS

* LEO PHARMA AS
 DOVONEX, CALCIPOTRIENE
 ENSTILAR, BETAMETHASONE DIPROPIONATE
 FINACEA, AZELAIC ACID
 PROTOPIC, TACROLIMUS
 TACLONEX, BETAMETHASONE DIPROPIONATE

LEXENPHARM

* LEXENPHARM SUZHOU LTD
 ALBUTEROL SULFATE, ALBUTEROL SULFATE
 ARFORMOTEROL TARTRATE, ARFORMOTEROL TARTRATE
 FORMOTEROL FUMARATE, FORMOTEROL FUMARATE

LEXICON PHARMS INC

* LEXICON PHARMACEUTICALS INC
 INPEFA, SOTAGLIFLOZIN

LGM PHARMA

* LGM PHARMA SOLUTIONS LLC
 BUTALBITAL AND ACETAMINOPHEN, ACETAMINOPHEN
 BUTALBITAL, ACETAMINOPHEN AND CAFFEINE, ACETAMINOPHEN
 BUTALBITAL, ACETAMINOPHEN, CAFFEINE AND CODEINE PHOSPHATE, ACETAMINOPHEN
 BUTALBITAL, ASPIRIN, CAFFEINE, AND CODEINE PHOSPHATE, ASPIRIN
 CHENODIOL, CHENODIOL
 GLYCOPYRROLATE, GLYCOPYRROLATE
 HYDROCODONE BITARTRATE AND ASPIRIN, ASPIRIN
 MECAMYLAMINE HYDROCHLORIDE, MECAMYLAMINE HYDROCHLORIDE
 NABUMETONE, NABUMETONE
 POLYETHYLENE GLYCOL 3350, POLYETHYLENE GLYCOL 3350 (OTC)
 URSODIOL, URSODIOL

LIEBEL-FLARSHEIM

* LIEBEL-FLARSHEIM CO LLC
 CONRAY, IOTHALAMATE MEGLUMINE
 CYSTO-CONRAY II, IOTHALAMATE MEGLUMINE
 MD-GASTROVIEW, DIATRIZOATE MEGLUMINE
 OPTIRAY 300, IOVERSOL
 OPTIRAY 320, IOVERSOL
 OPTIRAY 350, IOVERSOL
 SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, SODIUM CHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** L ******LIFE MOLECULAR**

* LIFE MOLECULAR IMAGING LTD
NEURACEQ, FLORBETABEN F-18

LINDE GAS EQUIP

* LINDE GAS AND EQUIPMENT INC
NOXIVENT, NITRIC OXIDE

LIVZON GRP

* LIVZON GROUP PHARMACEUTICAL FACTORY
CETRORELIX ACETATE, CETRORELIX ACETATE

LNHC

* LNHC INC
SITAVIG, ACYCLOVIR
ZELSUVMI, BERDAZIMER SODIUM

LNK

* LNK INTERNATIONAL INC
DOXYLAMINE SUCCINATE, DOXYLAMINE SUCCINATE (OTC)
IBUPROFEN, IBUPROFEN (OTC)
LOPERAMIDE HYDROCHLORIDE, LOPERAMIDE HYDROCHLORIDE (OTC)

LNK INTL INC

* LNK INTERNATIONAL INC
NAPROXEN SODIUM, NAPROXEN SODIUM (OTC)

LONG GROVE PHARMS

* LONG GROVE PHARMACEUTICALS LLC
AK-FLUOR 10%, FLUORESCEIN SODIUM
AK-FLUOR 25%, FLUORESCEIN SODIUM
CALCITRIOL, CALCITRIOL
CARBOPROST TROMETHAMINE, CARBOPROST TROMETHAMINE
DIAZEPAM, DIAZEPAM
EDARAVONE, EDARAVONE
ERIBULIN MESYLATE, ERIBULIN MESYLATE
METHADONE HYDROCHLORIDE, METHADONE HYDROCHLORIDE
NOREPINEPHRINE BITARTRATE IN 0.9% SODIUM CHLORIDE, NOREPINEPHRINE BITARTRATE
SODIUM BICARBONATE, SODIUM BICARBONATE
TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE
VASOPRESSIN IN SODIUM CHLORIDE 0.9%, VASOPRESSIN

LOREAL USA

* LOREAL USA PRODUCTS INC
ANTHELIOS 20, AVOBENZONE (OTC)
ANTHELIOS 40, AVOBENZONE (OTC)
ANTHELIOS SX, AVOBENZONE (OTC)
CAPITAL SOLEIL 15, AVOBENZONE (OTC)

LOTUS PHARM CO LTD

* LOTUS PHARMACEUTICAL CO LTD NANTOU PLANT
LENALIDOMIDE, LENALIDOMIDE

LOXO ONCOL

* LOXO ONCOLOGY INC
JAYPIRCA, PIRTOBRUTINIB

LOXO ONCOL ELI LILLY

* LOXO ONCOLOGY A WHOLLY OWNED SUB OF ELI LILLY AND COMPANY
RETEVMO, SELPERCATINIB
* LOXO ONCOLOGY INC A WHOLLY OWNED SUB OF ELI LILLY AND CO
RETEVMO, SELPERCATINIB

LUITPOLD

* LUITPOLD PHARMACEUTICALS INC
AMINOCAPROIC ACID, AMINOCAPROIC ACID

LUKARE MEDICAL LLC

* LUKARE MEDICAL LLC
ELLIOTTS B SOLUTION, CALCIUM CHLORIDE

LUMICELL

* LUMICELL INC
LUMISIGHT, PEGULICIANINE ACETATE

LUNDBECK NA LTD

* LUNDBECK NA LTD
NORTHERA, DROXIDOPA

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** L ******LUNDBECK PHARMS LLC**

* LUNDBECK PHARMACEUTICALS LLC
ONFI, CLOBAZAM
SABRIL, VIGABATRIN

LUOXIN AUROVITAS

* LUOXIN AUROVITAS PHARMA CHENGDU CO LTD
ALBUTEROL SULFATE, ALBUTEROL SULFATE
IPRATROPIUM BROMIDE, IPRATROPIUM BROMIDE
LEVALBUTEROL HYDROCHLORIDE, LEVALBUTEROL HYDROCHLORIDE

LUPIN

* LUPIN INC
ALBUTEROL SULFATE, ALBUTEROL SULFATE
ANTARA (MICRONIZED), FENOFIBRATE
ARFORMOTEROL TARTRATE, ARFORMOTEROL TARTRATE
ARIPIPIRAZOLE, ARIPIPIRAZOLE
BROVANA, ARFORMOTEROL TARTRATE
BUDESONIDE, BUDESONIDE
CHLORPROMAZINE HYDROCHLORIDE, CHLORPROMAZINE HYDROCHLORIDE
CYANOCOBALAMIN, CYANOCOBALAMIN
DESOXIMETASONE, DESOXIMETASONE
DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE
DIMETHYL FUMARATE, DIMETHYL FUMARATE
DOXORUBICIN HYDROCHLORIDE (LIPOSOMAL), DOXORUBICIN HYDROCHLORIDE
DOXYCYCLINE, DOXYCYCLINE
DRONEDARONE HYDROCHLORIDE, DRONEDARONE HYDROCHLORIDE
FLUOCINOLONE ACETONIDE, FLUOCINOLONE ACETONIDE
FORMOTEROL FUMARATE, FORMOTEROL FUMARATE
LEVOTHYROXINE SODIUM, LEVOTHYROXINE SODIUM **
MIBELAS 24 FE, ETHINYL ESTRADIOL
MIDOSTAURIN, MIDOSTAURIN
NALOXONE HYDROCHLORIDE AND PENTAZOCINE HYDROCHLORIDE, NALOXONE HYDROCHLORIDE
NYSTATIN, NYSTATIN
ORPHENADRINE CITRATE, ORPHENADRINE CITRATE
OSELTAMIVIR PHOSPHATE, OSELTAMIVIR PHOSPHATE
ROSUVASTATIN CALCIUM, ROSUVASTATIN CALCIUM
TESTOSTERONE, TESTOSTERONE
TIOTROPIUM BROMIDE, TIOTROPIUM BROMIDE
TOBRAMYCIN, TOBRAMYCIN
TRIMETHOBENZAMIDE HYDROCHLORIDE, TRIMETHOBENZAMIDE HYDROCHLORIDE
XOPENEX HFA, LEVALBUTEROL TARTRATE

* LUPIN LTD

AMLODIPINE BESYLATE AND VALSARTAN, AMLODIPINE BESYLATE
AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
CARVEDILOL, CARVEDILOL
CEFADROXIL, CEFADROXIL/CEFADROXIL HEMIHYDRATE
CEFDINIR, CEFDINIR
CEFPROZIL, CEFPROZIL
CEFUROXIME AXETIL, CEFUROXIME AXETIL
CEPHALEXIN, CEPHALEXIN
DIVALPROEX SODIUM, DIVALPROEX SODIUM
ETHAMBUTOL HYDROCHLORIDE, ETHAMBUTOL HYDROCHLORIDE
LEVETIRACETAM, LEVETIRACETAM
LEVOFLOXACIN, LEVOFLOXACIN
LISINAPRIL AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
LISINAPRIL, LISINAPRIL
LOVASTATIN, LOVASTATIN
QUINAPRIL HYDROCHLORIDE, QUINAPRIL HYDROCHLORIDE
RAMIPRIL, RAMIPRIL
SERTRALINE HYDROCHLORIDE, SERTRALINE HYDROCHLORIDE
SIMVASTATIN, SIMVASTATIN
TRANDOLAPRIL, TRANDOLAPRIL

LUPIN LTD

* LUPIN LIMITED
LEVETIRACETAM, LEVETIRACETAM
LEVONORGESTREL AND ETHINYL ESTRADIOL AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** L **

* LUPIN LTD

ABACAVIR SULFATE, LAMIVUDINE AND ZIDOVUDINE, ABACAVIR SULFATE
 AMABELZ, ESTRADIOL
 AMLODIPINE BESYLATE, VALSARTAN AND HYDROCHLOROTHIAZIDE, AMLODIPINE BESYLATE
 ARMODAFINIL, ARMODAFINIL
 ATOVAQUONE, ATOVAQUONE
 AZITHROMYCIN, AZITHROMYCIN
 BEKYREE, DESOGESTREL
 BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
 BIMATOPROST, BIMATOPROST
 BLISOVI 24 FE, ETHINYL ESTRADIOL
 BLISOVI FE 1.5/30, ETHINYL ESTRADIOL
 BLISOVI FE 1/20, ETHINYL ESTRADIOL
 BRIMONIDINE TARTRATE, BRIMONIDINE TARTRATE
 BROMFENAC SODIUM, BROMFENAC SODIUM
 BUMETANIDE, BUMETANIDE
 BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
 CALCIUM ACETATE, CALCIUM ACETATE
 CELECOXIB, CELECOXIB
 CLOBAZAM, CLOBAZAM
 CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
 CLOMIPRAMINE HYDROCHLORIDE, CLOMIPRAMINE HYDROCHLORIDE
 DARUNAVIR, DARUNAVIR
 DAYSEE, ETHINYL ESTRADIOL
 DECITABINE, DECITABINE
 DESVENLAFAXINE SUCCINATE, DESVENLAFAXINE SUCCINATE
 DICLOFENAC SODIUM, DICLOFENAC SODIUM
 DIVALPROEX SODIUM, DIVALPROEX SODIUM
 DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
 DOXERCALCIFEROL, DOXERCALCIFEROL
 DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
 DOXYCYCLINE, DOXYCYCLINE
 DROSPIRENONE AND ETHINYL ESTRADIOL, DROSPIRENONE
 DROSPIRENONE, ETHINYL ESTRADIOL AND LEVOMEFOLATE CALCIUM, DROSPIRENONE
 DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE
 EMTRICITABINE AND TENOFOVIR ALAFENAMIDE FUMARATE, EMTRICITABINE
 ENSKYCE, DESOGESTREL
 ESZOPICLONE, ESZOPICLONE
 ETHACRYNIC ACID, ETHACRYNIC ACID
 FALLBACK SOLO, LEVONORGESTREL (OTC)
 FAMOTIDINE, FAMOTIDINE
 FENOFIBRATE, FENOFIBRATE
 FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 FOSAPREPITANT DIMEGLUMINE, FOSAPREPITANT DIMEGLUMINE
 FYAVOLV, ETHINYL ESTRADIOL
 GANIRELIX ACETATE, GANIRELIX ACETATE
 GATIFLOXACIN, GATIFLOXACIN
 HYDROCORTISONE BUTYRATE, HYDROCORTISONE BUTYRATE
 HYDROCORTISONE VALERATE, HYDROCORTISONE VALERATE
 IMIPRAMINE PAMOATE, IMIPRAMINE PAMOATE
 IRBESARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 IRBESARTAN, IRBESARTAN
 JENCYCLA, NORETHINDRONE
 KAITLIB FE, ETHINYL ESTRADIOL
 KETOROLAC TROMETHAMINE AND PHENYLEPHRINE HYDROCHLORIDE, KETOROLAC TROMETHAMINE
 KURVELO, ETHINYL ESTRADIOL
 LAMIVUDINE AND ZIDOVUDINE, LAMIVUDINE
 LAMIVUDINE, LAMIVUDINE
 LAMOTRIGINE, LAMOTRIGINE
 LEFLUNOMIDE, LEFLUNOMIDE
 LEVETIRACETAM, LEVETIRACETAM
 LEVONORGESTREL AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
 LORAZEPAM, LORAZEPAM
 LOSARTAN POTASSIUM AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 LOSARTAN POTASSIUM, LOSARTAN POTASSIUM

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** L ******* LUPIN LTD**

LOTE Prednol Etabonate, LOTE Prednol Etabonate
 Lurasidone Hydrochloride, Lurasidone Hydrochloride
 Mefenamic Acid, Mefenamic Acid
 Meloxicam, Meloxicam
 Memantine Hydrochloride, Memantine Hydrochloride
 Metformin Hydrochloride, Metformin Hydrochloride
 Metronidazole, Metronidazole
 MinzoYA, Ethinyl Estradiol
 Mirabegron, Mirabegron
 Moxifloxacin Hydrochloride, Moxifloxacin Hydrochloride
 Nikki, Drospirenone
 Norethindrone and Ethinyl Estradiol and Ferrous Fumarate, Ethinyl Estradiol
 Norethindrone, Norethindrone
 Norgestimate and Ethinyl Estradiol, Ethinyl Estradiol
 Nystatin and Triamcinolone Acetonide, Nystatin
 Paliperidone, Paliperidone
 Paroxetine Hydrochloride, Paroxetine Hydrochloride
 Penicillamine, Penicillamine
 Pitavastatin Calcium, Pitavastatin Calcium
 Potassium Chloride, Potassium Chloride
 Prednisolone Acetate, Prednisolone Acetate
 Quetiapine Fumarate, Quetiapine Fumarate
 Quinine Sulfate, Quinine Sulfate
 Rifabutin, Rifabutin
 Rufinamide, Rufinamide
 Sevelamer Hydrochloride, Sevelamer Hydrochloride
 Sildenafil Citrate, Sildenafil Citrate
 Silodosin, Silodosin
 Suprax, Cefixime
 Tadalafil, Tadalafil
 Tavaborole, Tavaborole
 Telmisartan and Amlodipine, Amlodipine Besylate
 Telmisartan and Hydrochlorothiazide, Hydrochlorothiazide
 Tenofovir Alafenamide, Tenofovir Alafenamide Fumarate
 Testosterone, Testosterone
 Topiramate, Topiramate
 Tramadol Hydrochloride, Tramadol Hydrochloride
 Turqoz, Ethinyl Estradiol
 Tydemy, Drospirenone
 Valbenazine Tosylate, Valbenazine Tosylate
 Valsartan and Hydrochlorothiazide, Hydrochlorothiazide
 Valsartan, Valsartan
 Vancomycin Hydrochloride, Vancomycin Hydrochloride
 Varenicline Tartrate, Varenicline Tartrate
 Vyfemla, Ethinyl Estradiol
 Zolpidem Tartrate, Zolpidem Tartrate

LUPIN PHARMS*** LUPIN PHARMACEUTICALS INC**

Amlodipine Besylate and Benazepril Hydrochloride, Amlodipine Besylate
 Desloratadine, Desloratadine
 Diclofenac Sodium, Diclofenac Sodium
 Droxidopa, Droxidopa
 Meloxicam, Meloxicam
 Norgestimate and Ethinyl Estradiol, Ethinyl Estradiol
 Rifampin, Rifampin
 Suprax, Cefixime
 Ziprasidone Hydrochloride, Ziprasidone Hydrochloride

LUYE INNOMIND PHARMA*** LUYE INNOMIND PHARMA (SHIJIAZHUANG) CO LTD**

Erzofri, Paliperidone Palmitate

LYNE*** LYNE LABORATORIES INC**

Chlorhexidine Gluconate, Chlorhexidine Gluconate
 Nystatin, Nystatin

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** L ******PERRIGO**

* L PERRIGO CO

ACETAMINOPHEN, ACETAMINOPHEN (OTC)
 ACETAMINOPHEN, ASPIRIN AND CAFFEINE, ACETAMINOPHEN (OTC)
 CHILDREN'S IBUPROFEN, IBUPROFEN (OTC)
 DOXYLAMINE SUCCINATE, DOXYLAMINE SUCCINATE (OTC)
 FAMOTIDINE, FAMOTIDINE (OTC)
 IBUPROFEN, IBUPROFEN (OTC)
 LORATADINE, LORATADINE (OTC)
 MICONAZOLE 3 COMBINATION PACK, MICONAZOLE NITRATE (OTC)
 MICONAZOLE NITRATE, MICONAZOLE NITRATE (OTC)
 MINOXIDIL EXTRA STRENGTH (FOR MEN), MINOXIDIL (OTC)
 NAPROXEN SODIUM AND PSEUDOEPHEDRINE HYDROCHLORIDE, NAPROXEN SODIUM (OTC)
 NAPROXEN SODIUM, NAPROXEN SODIUM (OTC)
 TIOCONAZOLE, TIOCONAZOLE (OTC)

**** M ******MA GENERAL HOSP**

* MASSACHUSETTS GENERAL HOSP

FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18

MACLEODS PHARMS LTD

* MACLEODS PHARMACEUTICALS LTD

ABACAVIR SULFATE AND LAMIVUDINE, ABACAVIR SULFATE
 ACYCLOVIR, ACYCLOVIR
 AMLODIPINE AND OLMESARTAN MEDOXOMIL, AMLODIPINE BESYLATE
 AMLODIPINE BESYLATE AND VALSARTAN, AMLODIPINE BESYLATE
 AMLODIPINE BESYLATE, VALSARTAN AND HYDROCHLOROTHIAZIDE, AMLODIPINE BESYLATE
 APREMILAST, APREMILAST
 ARIPIPIRAZOLE, ARIPIPIRAZOLE
 ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM
 CANDESARTAN CILEXETIL AND HYDROCHLOROTHIAZIDE, CANDESARTAN CILEXETIL
 CANDESARTAN CILEXETIL, CANDESARTAN CILEXETIL
 CELECOXIB, CELECOXIB
 CEVIMELINE HYDROCHLORIDE, CEVIMELINE HYDROCHLORIDE
 CHOLESTYRAMINE LIGHT, CHOLESTYRAMINE
 CHOLESTYRAMINE, CHOLESTYRAMINE
 CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
 CLOPIDOGREL BISULFATE, CLOPIDOGREL BISULFATE
 CYCLOBENZAPRINE HYDROCHLORIDE, CYCLOBENZAPRINE HYDROCHLORIDE
 DARIFENACIN, DARIFENACIN HYDROBROMIDE
 DIMETHYL FUMARATE, DIMETHYL FUMARATE
 DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
 DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE
 EFAVIRENZ, EFAVIRENZ
 EFAVIRENZ, EMTRICITABINE, AND TENOFOVIR DISOPROXIL FUMARATE, EFAVIRENZ
 EMTRICITABINE AND TENOFOVIR DISOPROXIL FUMARATE, EMTRICITABINE
 ENTACAPONE, ENTACAPONE
 ESCITALOPRAM OXALATE, ESCITALOPRAM OXALATE
 ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM
 ESZOPICLONE, ESZOPICLONE
 EZETIMIBE, EZETIMIBE
 FAMCICLOVIR, FAMCICLOVIR
 FEBUXOSTAT, FEBUXOSTAT
 FENOFIBRATE, FENOFIBRATE
 FLUOCINONIDE, FLUOCINONIDE
 HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
 IBANDRONATE SODIUM, IBANDRONATE SODIUM
 IRBESARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 IRBESARTAN, IRBESARTAN
 LACOSAMIDE, LACOSAMIDE
 LAMIVUDINE AND ZIDOVUDINE, LAMIVUDINE
 LAMIVUDINE, LAMIVUDINE
 LANSOPRAZOLE, LANSOPRAZOLE
 LEVOCETIRIZINE DIHYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE
 LEVOFLOXACIN, LEVOFLOXACIN

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** M ****

* MACLEODS PHARMACEUTICALS LTD
 LEVOTHYROXINE SODIUM, LEVOTHYROXINE SODIUM **
 LIDOCAINE, LIDOCAINE
 LINEZOLID, LINEZOLID
 LOPINAVIR AND RITONAVIR, LOPINAVIR
 LOSARTAN POTASSIUM AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 LOSARTAN POTASSIUM, LOSARTAN POTASSIUM
 LURASIDONE HYDROCHLORIDE, LURASIDONE HYDROCHLORIDE
 MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 METHIMAZOLE, METHIMAZOLE
 MONTELUKAST SODIUM, MONTELUKAST SODIUM
 NEVIRAPINE, NEVIRAPINE
 NIACIN, NIACIN
 NYSTATIN AND TRIAMCINOLONE ACETONIDE, NYSTATIN
 NYSTATIN, NYSTATIN
 OLANZAPINE, OLANZAPINE
 OLMESARTAN MEDOXOMIL AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 OLMESARTAN MEDOXOMIL, OLMESARTAN MEDOXOMIL
 OSELTAMIVIR PHOSPHATE, OSELTAMIVIR PHOSPHATE
 PERPHENAZINE, PERPHENAZINE
 PIOGLITAZONE HYDROCHLORIDE AND METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 PIOGLITAZONE HYDROCHLORIDE, PIOGLITAZONE HYDROCHLORIDE
 PIRFENIDONE, PIRFENIDONE
 POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
 PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE
 PREGABALIN, PREGABALIN
 PROPYLTHIOURACIL, PROPYLTHIOURACIL
 PYRAZINAMIDE, PYRAZINAMIDE
 QUETIAPINE FUMARATE, QUETIAPINE FUMARATE
 RASAGILINE MESYLATE, RASAGILINE MESYLATE
 REPAGLINIDE, REPAGLINIDE
 RISEDRONATE SODIUM, RISEDRONATE SODIUM
 RIVASTIGMINE TARTRATE, RIVASTIGMINE TARTRATE
 RIZATRIPTAN BENZOATE, RIZATRIPTAN BENZOATE
 ROSUVASTATIN CALCIUM, ROSUVASTATIN CALCIUM
 SACUBITRIL AND VALSARTAN, SACUBITRIL
 SEVELAMER CARBONATE, SEVELAMER CARBONATE
 SEVELAMER HYDROCHLORIDE, SEVELAMER HYDROCHLORIDE
 SILDENAFIL CITRATE, SILDENAFIL CITRATE
 SILODOSIN, SILODOSIN
 SOLIFENACIN SUCCINATE, SOLIFENACIN SUCCINATE
 TADALAFIL, TADALAFIL
 TAMSULOSIN HYDROCHLORIDE, TAMSULOSIN HYDROCHLORIDE
 TENOFOVIR DISOPROXIL FUMARATE, TENOFOVIR DISOPROXIL FUMARATE
 TOLTERODINE TARTRATE, TOLTERODINE TARTRATE
 TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE
 TROSPIUM CHLORIDE, TROSPIUM CHLORIDE
 VALSARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 VALSARTAN, VALSARTAN
 VARDENAFIL HYDROCHLORIDE, VARDENAFIL HYDROCHLORIDE
 VARENICLINE TARTRATE, VARENICLINE TARTRATE
 VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE
 ZIPRASIDONE HYDROCHLORIDE, ZIPRASIDONE HYDROCHLORIDE

MADRIGAL

* MADRIGAL PHARMACEUTICALS INC
 REZDIFFRA, RESMETIROM

MAIA PHARMS INC

* MAIA PHARMACEUTICALS INC
 ANGIOMAX RTU, BIVALIRUDIN
 BACLOFEN, BACLOFEN
 DAPTOMYCIN, DAPTOMYCIN
 LEVOTHYROXINE SODIUM, LEVOTHYROXINE SODIUM
 SINCALIDE, SINCALIDE
 SODIUM PHENYLACETATE AND SODIUM BENZOATE, SODIUM BENZOATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** M ******MAINPOINTE**

* MAINPOINTE PHARMACEUTICALS LLC
TUXARIN ER, CHLORPHENIRAMINE MALEATE

MALLINCKRODT HOSP

* MALLINCKRODT HOSP PRODUCTS IP LTD
INOMAX, NITRIC OXIDE

MALLINCKRODT IRELAND

* MALLINCKRODT PHARMACEUTICALS IRELAND LTD
ACTHAR GEL (AUTOINJECTOR), CORTICOTROPIN
ACTHAR GEL, CORTICOTROPIN
TERLIVAZ, TERLIPRESSIN ACETATE

MANKIND PHARMA

* MANKIND PHARMA LTD
ACETAZOLAMIDE, ACETAZOLAMIDE
ALOSETRON HYDROCHLORIDE, ALOSETRON HYDROCHLORIDE
AMITRIPTYLINE HYDROCHLORIDE, AMITRIPTYLINE HYDROCHLORIDE
APREMILAST, APREMILAST
ARFORMOTEROL TARTRATE, ARFORMOTEROL TARTRATE
ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM
ATROPINE SULFATE, ATROPINE SULFATE
BACLOFEN, BACLOFEN
CHLORTHALIDONE, CHLORTHALIDONE
CLOMIPRAMINE HYDROCHLORIDE, CLOMIPRAMINE HYDROCHLORIDE
CYANOCOBALAMIN, CYANOCOBALAMIN
DOXEPIN HYDROCHLORIDE, DOXEPIN HYDROCHLORIDE
EPHEDRINE SULFATE, EPHEDRINE SULFATE
FAMOTIDINE, FAMOTIDINE
FENOFIBRATE, FENOFIBRATE
FORMOTEROL FUMARATE, FORMOTEROL FUMARATE
FUROSEMIDE, FUROSEMIDE
HALOPERIDOL DECANOATE, HALOPERIDOL DECANOATE
HALOPERIDOL, HALOPERIDOL
LEVALBUTEROL HYDROCHLORIDE, LEVALBUTEROL HYDROCHLORIDE
LIDOCAINE HYDROCHLORIDE, LIDOCAINE HYDROCHLORIDE
MIDODRINE HYDROCHLORIDE, MIDODRINE HYDROCHLORIDE
NALOXONE HYDROCHLORIDE, NALOXONE HYDROCHLORIDE
NEBIVOLOL HYDROCHLORIDE, NEBIVOLOL HYDROCHLORIDE
NITROFURANTOIN (MONOHYDRATE/MACROCRYSTALS), NITROFURANTOIN
NITROFURANTOIN, NITROFURANTOIN, MACROCRYSTALLINE
NITROGLYCERIN, NITROGLYCERIN
OFLOXACIN, OFLOXACIN
OMEGA-3-ACID ETHYL ESTERS, OMEGA-3-ACID ETHYL ESTERS
PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
PHENYLEPHRINE HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE
PRAZOSIN HYDROCHLORIDE, PRAZOSIN HYDROCHLORIDE
RANOLAZINE, RANOLAZINE
TELMISARTAN, TELMISARTAN
TIMOLOL MALEATE, TIMOLOL MALEATE
TOBRAMYCIN, TOBRAMYCIN
VARENICLINE TARTRATE, VARENICLINE TARTRATE
VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE

MARINUS

* MARINUS PHARMACEUTICALS INC
ZTALMY, GANAXOLONE

MARIUS PHARMS LLC

* MARIUS PHARMACEUTICALS LLC
KYZATREX, TESTOSTERONE UNDECANOATE

MARKSANS PHARMA

* MARKSANS PHARMA LTD
ACETAMINOPHEN AND IBUPROFEN, ACETAMINOPHEN (OTC)
ACETAMINOPHEN, ACETAMINOPHEN (OTC)
CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
CETIRIZINE HYDROCHLORIDE HIVES, CETIRIZINE HYDROCHLORIDE (OTC)
ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM (OTC)
FAMOTIDINE, FAMOTIDINE (OTC)

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** M ******* MARKSANS PHARMA LTD**

FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 GABAPENTIN, GABAPENTIN
 GUAIFENESIN, GUAIFENESIN (OTC)
 IBUPROFEN, IBUPROFEN
 IBUPROFEN, IBUPROFEN (OTC)
 LORATADINE, LORATADINE (OTC)
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 NAPROXEN SODIUM, NAPROXEN SODIUM (OTC)
 PARICALCITOL, PARICALCITOL

MAYNE PHARMA*** MAYNE PHARMA INTERNATIONAL PTY LTD**

DORYX MPC, DOXYCYCLINE HYCLATE
 DORYX, DOXYCYCLINE HYCLATE
 TOLSURA, ITRACONAZOLE

*** MAYNE PHARMA LLC**

ANNOVERA, ETHINYL ESTRADIOL
 BIJUVA, ESTRADIOL
 FABIOR, TAZAROTENE
 IMVEXXY, ESTRADIOL
 LEXETTE, HALOBETASOL PROPIONATE
 NEXTSTELLIS, DROSPIRENONE
 RHOFAD, OXYMETAZOLINE HYDROCHLORIDE
 SORILUX, CALCIPOTRIENE

MAYNE PHARMA COMMRCCL*** MAYNE PHARMA COMMERCIAL LLC**

SOLTAMOX, TAMOXIFEN CITRATE

MC2*** MC2 THERAPEUTICS LTD**

WYNZORA, BETAMETHASONE DIPROPIONATE

MCGUFF*** MCGUFF PHARMACEUTICALS INC**

ASCOR, ASCORBIC ACID

MCNEIL*** MCNEIL CONSUMER PRODUCTS CO DIV MCNEILAB INC**

IBUPROFEN, IBUPROFEN (OTC)

MCPRF*** MAYO CLINIC PET RADIOCHEMISTRY FACILITY**

AMMONIA N 13, AMMONIA N-13
 CHOLINE C-11, CHOLINE C-11
 FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18
 SODIUM FLUORIDE F-18, SODIUM FLUORIDE F-18

MDD US*** MDD US OPERATIONS LLC**

APOKYN, APOMORPHINE HYDROCHLORIDE
 XADAGO, SAFINAMIDE MESYLATE

MDGH*** MEDICINES DEVELOPMENT FOR GLOBAL HEALTH**

MOXIDECTIN, MOXIDECTIN

MEDEFIL INC*** MEDEFIL INC**

ATROPINE SULFATE, ATROPINE SULFATE
 CALCIUM CHLORIDE 10%, CALCIUM CHLORIDE
 SODIUM CHLORIDE 0.9%, SODIUM CHLORIDE
 STERILE WATER FOR INJECTION, STERILE WATER FOR INJECTION

MEDEXUS*** MEDEXUS PHARMA INC**

RASUVO, METHOTREXATE

MEDI-PHYSICS*** MEDI-PHYSICS INC DBA GE HEALTHCARE**

MYOVIEV 30ML, TECHNETIUM TC-99M TETROFOSMIN KIT
 MYOVIEV, TECHNETIUM TC-99M TETROFOSMIN KIT

MEDI-RADIOPHARMA*** MEDI-RADIOPHARMA LTD**

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** M ****

* MEDI-RADIOPHARMA LTD
TECHNETIUM TC99M MERTIATIDE KIT, TECHNETIUM TC-99M MERTIATIDE KIT

MEDICINES360

* MEDICINES360
LILETTA, LEVONORGESTREL

MEDICURE

* MEDICURE INTERNATIONAL INC
AGGRASTAT, TIROFIBAN HYDROCHLORIDE
ZYPITAMAG, PITAVASTATIN MAGNESIUM

MEDIMETRIKS PHARMS

* MEDIMETRIKS PHARMACEUTICALS INC
BETAXOLOL HYDROCHLORIDE, BETAXOLOL HYDROCHLORIDE
LOPROX, CICLOPIROX
NEO-SYNALAR, FLUOCINOLONE ACETONIDE
SYNALAR, FLUOCINOLONE ACETONIDE

MEDLEY PHARMS

* MEDLEY PHARMACEUTICALS LTD
LACOSAMIDE, LACOSAMIDE
NYSTATIN, NYSTATIN
VARENICLINE TARTRATE, VARENICLINE TARTRATE

MEDLINE INDUSTRIES

* MEDLINE INDUSTRIES INC
READYPREP CHG, CHLORHEXIDINE GLUCONATE (OTC)

MEDETECH PRODUCTS

* MEDETECH PRODUCTS INC
MONISTAT 1 COMBINATION PACK, MICONAZOLE NITRATE (OTC)
MONISTAT 3 COMBINATION PACK (PREFILLED), MICONAZOLE NITRATE (OTC)
MONISTAT 3 COMBINATION PACK, MICONAZOLE NITRATE (OTC)
MONISTAT 3, MICONAZOLE NITRATE
MONISTAT 3, MICONAZOLE NITRATE (OTC)
MONISTAT 7 COMBINATION PACK, MICONAZOLE NITRATE (OTC)
MONISTAT 7, MICONAZOLE NITRATE (OTC)
NIX, PERMETHRIN (OTC)
TAGAMET HB, CIMETIDINE (OTC)

MEDUNIK

* MEDUNIK CANADA INC
NITISINONE, NITISINONE
PHEBURANE, SODIUM PHENYLBUTYRATE

MEITHEAL

* MEITHEAL PHARMACEUTICALS INC
ADENOSINE, ADENOSINE
ALPROSTADIL, ALPROSTADIL
AMIKACIN SULFATE, AMIKACIN SULFATE
ATRACURIUM BESYLATE PRESERVATIVE FREE, ATRACURIUM BESYLATE
ATRACURIUM BESYLATE, ATRACURIUM BESYLATE
AZACITIDINE, AZACITIDINE
BENDAMUSTINE HYDROCHLORIDE, BENDAMUSTINE HYDROCHLORIDE
BIVALIRUDIN, BIVALIRUDIN
BLEOMYCIN SULFATE, BLEOMYCIN SULFATE
BORTEZOMIB, BORTEZOMIB
BUPIVACAINE HYDROCHLORIDE, BUPIVACAINE HYDROCHLORIDE
BUSULFAN, BUSULFAN
CARMUSTINE, CARMUSTINE
CISATRACURIUM BESYLATE, CISATRACURIUM BESYLATE
CLOFARABINE, CLOFARABINE
CYTARABINE, CYTARABINE
DACARBAZINE, DACARBAZINE
DACTINOMYCIN, DACTINOMYCIN
DAPTOMYCIN, DAPTOMYCIN
DAUNORUBICIN HYDROCHLORIDE, DAUNORUBICIN HYDROCHLORIDE
DECITABINE, DECITABINE
DESMOPRESSIN ACETATE, DESMOPRESSIN ACETATE
DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE
DOCETAXEL, DOCETAXEL
DOXERCALCIFEROL, DOXERCALCIFEROL

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** M ****

* MEITHEAL PHARMACEUTICALS INC
 EPOPROSTENOL SODIUM, EPOPROSTENOL SODIUM
 ETOPOSIDE, ETOPOSIDE
 FUROSEMIDE, FUROSEMIDE
 GANIRELIX ACETATE, GANIRELIX ACETATE
 GEMCITABINE HYDROCHLORIDE, GEMCITABINE HYDROCHLORIDE
 GLYCOPYRROLATE, GLYCOPYRROLATE
 HALOPERIDOL DECANOATE, HALOPERIDOL DECANOATE
 IDARUBICIN HYDROCHLORIDE, IDARUBICIN HYDROCHLORIDE
 IFOSFAMIDE, IFOSFAMIDE
 ISOSULFAN BLUE, ISOSULFAN BLUE
 LEUCOVORIN CALCIUM, LEUCOVORIN CALCIUM
 LEUPROLIDE ACETATE, LEUPROLIDE ACETATE
 LEVOLEUCOVORIN CALCIUM, LEVOLEUCOVORIN CALCIUM
 MELPHALAN HYDROCHLORIDE, MELPHALAN HYDROCHLORIDE
 MICAFUNGIN SODIUM, MICAFUNGIN SODIUM
 MILRINONE LACTATE, MILRINONE LACTATE
 MITOMYCIN, MITOMYCIN
 MITOXANTRONE HYDROCHLORIDE, MITOXANTRONE HYDROCHLORIDE
 MYCOPHENOLATE MOFETIL HYDROCHLORIDE, MYCOPHENOLATE MOFETIL HYDROCHLORIDE
 NELARABINE, NELARABINE
 NEOSTIGMINE METHYLSULFATE, NEOSTIGMINE METHYLSULFATE
 NOREPINEPHRINE BITARTRATE, NOREPINEPHRINE BITARTRATE
 OCTREOTIDE ACETATE, OCTREOTIDE ACETATE
 OXALIPLATIN, OXALIPLATIN
 PALONOSETRON HYDROCHLORIDE, PALONOSETRON HYDROCHLORIDE
 PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
 PEMETREXED DISODIUM, PEMETREXED DISODIUM
 PHENYLEPHRINE HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE
 PLERIXAFOR, PLERIXAFOR
 REGADENOSON, REGADENOSON
 ROCURONIUM BROMIDE, ROCURONIUM BROMIDE
 SUCCINYLCHOLINE CHLORIDE, SUCCINYLCHOLINE CHLORIDE
 THIOTEPA, THIOTEPA
 TIGECYCLINE, TIGECYCLINE
 VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE
 VECURONIUM BROMIDE, VECURONIUM BROMIDE
 VORICONAZOLE, VORICONAZOLE
 ZOLEDRONIC ACID, ZOLEDRONIC ACID

MELINTA

* MELINTA SUBSIDIARY CORP
 BAXDELA, DELAFLOXACIN MEGLUMINE

MELINTA THERAP

* MELINTA THERAPEUTICS LLC
 KIMYRSA, ORITAVANCIN DIPHOSPHATE
 ORBACTIV, ORITAVANCIN DIPHOSPHATE

MEM SLOAN-KETTERING

* MEMORIAL SLOAN-KETTERING CANCER CENTER
 FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18

MERCK

* MERCK AND CO INC
 EMEND, APREPITANT
 PRIMAXIN, CILASTATIN SODIUM

MERCK AND CO INC

* MERCK AND CO INC
 EMEND, FOSAPREPITANT DIMEGLUMINE

MERCK SHARP DOHME

* MERCK SHARP AND DOHME CORP
 BELSOMRA, SUVOREXANT
 JANUVIA, SITAGLIPTIN PHOSPHATE
 NOXAFIL, POSACONAZOLE
 PREVYMIS, LETERMIVIR
 STROMEKTOL, IVERMECTIN
 TEMODAR, TEMOZOLOMIDE

* MERCK SHARP AND DOHME CORP A SUB OF MERCK AND CO INC

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** M ****

* MERCK SHARP AND DOHME CORP A SUB OF MERCK AND CO INC
WELIREG, BELZUTIFAN

MERIDIAN BIOSCIENCE

* MERIDIAN BIOSCIENCE ISRAEL LTD
IDKIT:HP, CITRIC ACID

MERRO PHARM USA

* MERRO PHARMACEUTICAL USA INC
PHENTERMINE HYDROCHLORIDE, PHENTERMINE HYDROCHLORIDE
TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE

MERZ PHARMS

* MERZ PHARMACEUTICALS LLC
CUVPOSA, GLYCOPYRROLATE
* MERZ PHARMACEUTICALS LLC A SUB OF MERZ THERAPEUTICS GMBH
AMPYRA, DALFAMPRIDINE
INBRIJA, LEVODOPA

METACEL PHARMS LLC

* METACEL PHARMACEUTICALS LLC
OZOBAX DS, BACLOFEN

METHAPHARM

* METHAPHARM INC
PROVOCHOLINE, METHACHOLINE CHLORIDE

METUCHEN PHARMS

* METUCHEN PHARMACEUTICALS LLC
STENDRA, AVANAFIL

MICRO LABS

* MICRO LABS LTD
AMLODIPINE AND OLMESARTAN MEDOXOMIL, AMLODIPINE BESYLATE
AMOXICILLIN AND CLAVULANATE POTASSIUM, AMOXICILLIN
AMOXICILLIN, AMOXICILLIN
ASPIRIN AND DIPYRIDAMOLE, ASPIRIN
BACLOFEN, BACLOFEN
BIMATOPROST, BIMATOPROST
BRIMONIDINE TARTRATE, BRIMONIDINE TARTRATE
BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
CAFFEINE CITRATE, CAFFEINE CITRATE
CARBOPROST TROMETHAMINE, CARBOPROST TROMETHAMINE
CELECOXIB, CELECOXIB
CHLORDIAZEPOXIDE AND AMITRIPTYLINE HYDROCHLORIDE, AMITRIPTYLINE HYDROCHLORIDE
CHLORDIAZEPOXIDE HYDROCHLORIDE AND CLIDINIUM BROMIDE, CHLORDIAZEPOXIDE HYDROCHLORIDE
CLINDAMYCIN HYDROCHLORIDE, CLINDAMYCIN HYDROCHLORIDE
CLOBAZAM, CLOBAZAM
CLOMIPRAMINE HYDROCHLORIDE, CLOMIPRAMINE HYDROCHLORIDE
CROMOLYN SODIUM, CROMOLYN SODIUM
DALFAMPRIDINE, DALFAMPRIDINE
DEXAMETHASONE SODIUM PHOSPHATE, DEXAMETHASONE SODIUM PHOSPHATE
DICLOFENAC SODIUM AND MISOPROSTOL, DICLOFENAC SODIUM
DIPHENHYDRAMINE HYDROCHLORIDE, DIPHENHYDRAMINE HYDROCHLORIDE
DORZOLAMIDE HYDROCHLORIDE AND TIMOLOL MALEATE, DORZOLAMIDE HYDROCHLORIDE
DORZOLAMIDE HYDROCHLORIDE, DORZOLAMIDE HYDROCHLORIDE
DOXEPIN HYDROCHLORIDE, DOXEPIN HYDROCHLORIDE
ERYTHRA-DERM, ERYTHROMYCIN
FAMOTIDINE, FAMOTIDINE
FENOFIBRIC ACID, CHOLINE FENOFIBRATE
FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
FORMOTEROL FUMARATE, FORMOTEROL FUMARATE
GLIMEPIRIDE, GLIMEPIRIDE
ISOPROTERENOL HYDROCHLORIDE, ISOPROTERENOL HYDROCHLORIDE
LEVETIRACETAM, LEVETIRACETAM
LEVOCETIRIZINE DIHYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE (OTC)
LINCOMYCIN HYDROCHLORIDE, LINCOMYCIN HYDROCHLORIDE
LOSARTAN POTASSIUM, LOSARTAN POTASSIUM
MEFENAMIC ACID, MEFENAMIC ACID
METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
METHENAMINE HIPPURATE, METHENAMINE HIPPURATE
METOLAZONE, METOLAZONE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** M ******* MICRO LABS LTD**

MIDAZOLAM HYDROCHLORIDE, MIDAZOLAM HYDROCHLORIDE
 MISOPROSTOL, MISOPROSTOL
 NICARDIPINE HYDROCHLORIDE, NICARDIPINE HYDROCHLORIDE
 OLMESARTAN MEDOXOMIL, OLMESARTAN MEDOXOMIL
 PIRFENIDONE, PIRFENIDONE
 PIROXICAM, PIROXICAM
 RAMELTEON, RAMELTEON
 RANOLAZINE, RANOLAZINE
 RASAGILINE MESYLATE, RASAGILINE MESYLATE
 ROFLUMILAST, ROFLUMILAST
 RUFINAMIDE, RUFINAMIDE
 SEVELAMER CARBONATE, SEVELAMER CARBONATE
 SIMVASTATIN, SIMVASTATIN
 SODIUM NITROPRUSSIDE, SODIUM NITROPRUSSIDE
 SUCCINYLCHOLINE CHLORIDE, SUCCINYLCHOLINE CHLORIDE
 TAFLUPROST, TAFLUPROST
 TELMISARTAN, TELMISARTAN
 TIMOLOL MALEATE, TIMOLOL MALEATE
 TOBRAMYCIN, TOBRAMYCIN
 TRANEXAMIC ACID, TRANEXAMIC ACID
 TRAVOPROST, TRAVOPROST
 TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE
 VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE

MICRO LABS LTD*** MICRO LABS LTD**

NEVIRAPINE, NEVIRAPINE

MICRO LABS LTD INDIA*** MICRO LABS LTD INDIA**

ACETAZOLAMIDE, ACETAZOLAMIDE
 AMOXICILLIN AND CLAVULANATE POTASSIUM, AMOXICILLIN
 ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM
 CROMOLYN SODIUM, CROMOLYN SODIUM
 KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
 LEVOCETIRIZINE DIHYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE
 LEVOFLOXACIN, LEVOFLOXACIN
 TRAMADOL HYDROCHLORIDE AND ACETAMINOPHEN, ACETAMINOPHEN

MIDWEST MEDCL*** MIDWEST MEDICAL ISOTOPES LLC CYCLOTRON DIV**

AMMONIA N 13, AMMONIA N-13
 SODIUM FLUORIDE F-18, SODIUM FLUORIDE F-18

MIKART*** MIKART LLC**

BUTALBITAL AND ACETAMINOPHEN, ACETAMINOPHEN
 BUTALBITAL, ACETAMINOPHEN AND CAFFEINE, ACETAMINOPHEN
 BUTAPAP, ACETAMINOPHEN
 CARBINOXAMINE MALEATE, CARBINOXAMINE MALEATE
 CHLORZOXAZONE, CHLORZOXAZONE
 ERGOTAMINE TARTRATE AND CAFFEINE, CAFFEINE
 ETHOSUXIMIDE, ETHOSUXIMIDE
 HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
 METHAZOLAMIDE, METHAZOLAMIDE
 METHOCARBAMOL, METHOCARBAMOL
 OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN
 TRIHEXYPHENIDYL HYDROCHLORIDE, TRIHEXYPHENIDYL HYDROCHLORIDE

MILLA PHARMS*** MILLA PHARMACEUTICALS INC**

DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE
 MAGNESIUM SULFATE IN PLASTIC CONTAINER, MAGNESIUM SULFATE
 PYRIDOSTIGMINE BROMIDE, PYRIDOSTIGMINE BROMIDE
 SODIUM ACETATE, SODIUM ACETATE
 TROMETHAMINE, TROMETHAMINE

MILLICENT*** MILLICENT PHARMA LTD**

INTRAROSA, PRASTERONE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** M ******MILLICENT PR**

* MILLICENT PUERTO RICO LLC
 FEMLYV, ETHINYL ESTRADIOL
 FEMRING, ESTRADIOL ACETATE

MIPS CRF

* MIPS CYCLOTRON AND RADIOCHEMISTRY FACILITY
 AMMONIA N 13, AMMONIA N-13
 FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18
 SODIUM FLUORIDE F-18, SODIUM FLUORIDE F-18

MIRUM

* MIRUM PHARMACEUTICALS INC
 CHOLBAM, CHOLIC ACID
 LIVMARLI, MARALIXIBAT CHLORIDE

MISEMER

* MISEMER PHARMACEUTICALS INC
 CHLORDIAZEPOXIDE HYDROCHLORIDE AND CLIDINIUM BROMIDE, CHLORDIAZEPOXIDE HYDROCHLORIDE
 KETOPROFEN, KETOPROFEN

MISSION PHARMA

* MISSION PHARMACAL CO
 LITHOSTAT, ACETOHYDROXAMIC ACID
 PREDNISOLONE SODIUM PHOSPHATE, PREDNISOLONE SODIUM PHOSPHATE
 TEXACORT, HYDROCORTISONE
 THIOLA, TIOPRONIN
 TINDAMAX, TINIDAZOLE
 UROCIT-K, POTASSIUM CITRATE

MISSION PHARMACAL

* MISSION PHARMACAL CO
 CARBINOXAMINE MALEATE, CARBINOXAMINE MALEATE
 GABAPENTIN, GABAPENTIN
 POTASSIUM IODIDE, POTASSIUM IODIDE (OTC)
 THIOLA EC, TIOPRONIN

MITSUBISHI TANABE

* MITSUBISHI TANABE PHARMA CORP
 RADICAVA ORS, EDARAVONE
 RADICAVA, EDARAVONE

MLV

* MLV PHARMA LLC
 CARVEDILOL, CARVEDILOL
 PYRIDOSTIGMINE BROMIDE, PYRIDOSTIGMINE BROMIDE
 ROPINIROLE HYDROCHLORIDE, ROPINIROLE HYDROCHLORIDE

MMT

* MERIDIAN MEDICAL TECHNOLOGIES LLC
 DUODOTE, ATROPINE
 SEIZALAM, MIDAZOLAM HYDROCHLORIDE

MOBIUS THERAP

* MOBIUS THERAPEUTICS LLC
 MITOSOL, MITOMYCIN

MOLNLYCKE HLTH

* MOLNLYCKE HEALTH CARE
 HIBICLENS, CHLORHEXIDINE GLUCONATE (OTC)
 HIBISTAT, CHLORHEXIDINE GLUCONATE (OTC)

MONARCH PHARMS

* MONARCH PHARMACEUTICALS LLC
 NEOSPORIN G.U. IRRIGANT, NEOMYCIN SULFATE
 NEOSPORIN, GRAMICIDIN
 SEPTRA DS, SULFAMETHOXAZOLE
 SEPTRA, SULFAMETHOXAZOLE
 VIROPTIC, TRIFLURIDINE

MONTEREY PHARMS LLC

* MONTEREY PHARMACEUTICALS LLC
 METHOCARBAMOL, METHOCARBAMOL

MOUNTAIN

* MOUNTAIN LLC
 CYPROHEPTADINE HYDROCHLORIDE, CYPROHEPTADINE HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** M ****

* MOUNTAIN LLC
 GRISEOFULVIN, ULTRAMICROSIZE, GRISEOFULVIN, ULTRAMICROSIZE
 METAXALONE, METAXALONE
 METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE

MSD

* MERCK SHARP AND DOHME LLC
 PREVYMIS, LETERMOVIR
 VERQUVO, VERICIGUAT

MSD MERCK CO

* MERCK SHARP AND DOHME CORP A SUB OF MERCK AND CO INC
 DELSTRIGO, DORAVIRINE
 EMEND, APREPITANT
 NOXAFIL POWDERMIX KIT, POSACONAZOLE
 PIFELTRO, DORAVIRINE
 RECARBRIO, CILASTATIN SODIUM

MSD SUB MERCK

* MERCK SHARP AND DOHME LLC A SUB OF MERCK AND CO INC
 BRIDION, SUGAMMADEX SODIUM
 INVANZ, ERTAPENEM SODIUM
 ISENTRESS HD, RALTEGRAVIR POTASSIUM
 ISENTRESS, RALTEGRAVIR POTASSIUM
 JANUMET XR, METFORMIN HYDROCHLORIDE
 JANUMET, METFORMIN HYDROCHLORIDE
 SEGLUROMET, ERTUGLIFLOZIN
 STEGLATRO, ERTUGLIFLOZIN
 STEGLUJAN, ERTUGLIFLOZIN
 ZEPATIER, ELBASVIR
 ZOLINZA, VORINOSTAT

MSN

* MSN LABORATORIES PRIVATE LTD
 ABIRATERONE ACETATE, ABIRATERONE ACETATE
 ACYCLOVIR, ACYCLOVIR
 ALBENDAZOLE, ALBENDAZOLE
 AMINOCAPROIC ACID, AMINOCAPROIC ACID
 APREMILAST, APREMILAST
 ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM
 BORTEZOMIB, BORTEZOMIB
 BUMETANIDE, BUMETANIDE
 CAPECITABINE, CAPECITABINE
 CARMUSTINE, CARMUSTINE
 CHLORPROMAZINE HYDROCHLORIDE, CHLORPROMAZINE HYDROCHLORIDE
 CLOBAZAM, CLOBAZAM
 CLOFARABINE, CLOFARABINE
 CLOPIDOGREL BISULFATE, CLOPIDOGREL BISULFATE
 DABIGATRAN ETEXILATE MESYLATE, DABIGATRAN ETEXILATE MESYLATE
 DARUNAVIR, DARUNAVIR
 DECITABINE, DECITABINE
 DEFERASIROX, DEFERASIROX
 DIMETHYL FUMARATE, DIMETHYL FUMARATE
 DOFETILIDE, DOFETILIDE
 DOXEPIN HYDROCHLORIDE, DOXEPIN HYDROCHLORIDE
 ERLOTINIB HYDROCHLORIDE, ERLOTINIB HYDROCHLORIDE
 FAMOTIDINE, FAMOTIDINE
 FEBUXOSTAT, FEBUXOSTAT
 FLUPHENAZINE DECANOATE, FLUPHENAZINE DECANOATE
 FLUPHENAZINE HYDROCHLORIDE, FLUPHENAZINE HYDROCHLORIDE
 FOSAPREPITANT DIMEGLUMINE, FOSAPREPITANT DIMEGLUMINE
 HALOPERIDOL, HALOPERIDOL
 ISOSULFAN BLUE, ISOSULFAN BLUE
 LACOSAMIDE, LACOSAMIDE
 LEVETIRACETAM, LEVETIRACETAM
 LISDEXAMFETAMINE DIMESYLATE, LISDEXAMFETAMINE DIMESYLATE
 LOSARTAN POTASSIUM, LOSARTAN POTASSIUM
 LURASIDONE HYDROCHLORIDE, LURASIDONE HYDROCHLORIDE
 MOXIFLOXACIN HYDROCHLORIDE, MOXIFLOXACIN HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** M ****

- * MSN LABORATORIES PRIVATE LTD
 NEBIVOLOL HYDROCHLORIDE, NEBIVOLOL HYDROCHLORIDE
 NELARABINE, NELARABINE
 OLMESARTAN MEDOXOMIL, OLMESARTAN MEDOXOMIL
 OSELTAMIVIR PHOSPHATE, OSELTAMIVIR PHOSPHATE
 PACLITAXEL, PACLITAXEL
 PIRFENIDONE, PIRFENIDONE
 PLERIXAFOR, PLERIXAFOR
 PRAZOSIN HYDROCHLORIDE, PRAZOSIN HYDROCHLORIDE
 PREGABALIN, PREGABALIN
 PYRIDOSTIGMINE BROMIDE, PYRIDOSTIGMINE BROMIDE
 RIOCIQUAT, RIOCIQUAT
 ROFLUMILAST, ROFLUMILAST
 ROSUVASTATIN CALCIUM, ROSUVASTATIN CALCIUM
 SACUBITRIL AND VALSARTAN, SACUBITRIL
 SAFINAMIDE MESYLATE, SAFINAMIDE MESYLATE
 SILDENAFIL CITRATE, SILDENAFIL CITRATE
 SILODOSIN, SILODOSIN
 SIROLIMUS, SIROLIMUS
 SOLIFENACIN SUCCINATE, SOLIFENACIN SUCCINATE
 TASIMELTEON, TASIMELTEON
 TERIFLUNOMIDE, TERIFLUNOMIDE
 THIOTEPA, THIOTEPA
 TIAGABINE HYDROCHLORIDE, TIAGABINE HYDROCHLORIDE
 TIPIRACIL HYDROCHLORIDE AND TRIFLURIDINE, TIPIRACIL HYDROCHLORIDE
 TOLVAPTAN, TOLVAPTAN
 TOREMIFENE CITRATE, TOREMIFENE CITRATE
 TRIENTINE HYDROCHLORIDE, TRIENTINE HYDROCHLORIDE
 VALGANCICLOVIR HYDROCHLORIDE, VALGANCICLOVIR HYDROCHLORIDE
 VIGABATRIN, VIGABATRIN
 ZIPRASIDONE MESYLATE, ZIPRASIDONE MESYLATE
- MSN LABS PVT LTD**
- * MSN LABORATORIES PRIVATE LTD
 AZACITIDINE, AZACITIDINE
 LACOSAMIDE, LACOSAMIDE
- MSN PHARMS INC**
- * MSN PHARMACEUTICALS INC
 DROXIDOPA, DROXIDOPA
- MUNDIPHARMA**
- * MUNDIPHARMA GMBH
 REZZAYO, REZAFUNGIN ACETATE
- MYCOVIA PHARMS**
- * MYCOVIA PHARMACEUTICALS INC
 VIVJOA, OTESECONAZOLE
- MYLAN**
- * MYLAN INSTITUTIONAL LLC A VIATRIS CO
 ALOPRIM, ALLOPURINOL SODIUM
- * MYLAN PHARMACEUTICALS
 METOPROLOL TARTRATE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
- * MYLAN PHARMACEUTICALS INC
 ABIRATERONE ACETATE, ABIRATERONE ACETATE
 ACAMPROSATE CALCIUM, ACAMPROSATE CALCIUM
 ACETAMINOPHEN, ACETAMINOPHEN
 ACITRETIN, ACITRETIN
 ALLOPURINOL, ALLOPURINOL
 ALMOTRIPTAN MALATE, ALMOTRIPTAN MALATE
 AMBRISENTAN, AMBRISENTAN
 AMLODIPINE BESYLATE AND ATORVASTATIN CALCIUM, AMLODIPINE BESYLATE
 AMLODIPINE BESYLATE AND VALSARTAN, AMLODIPINE BESYLATE
 ARIPIPRAZOLE, ARIPIPRAZOLE
 ATENOLOL, ATENOLOL
 ATOVAQUONE AND PROGUANIL HYDROCHLORIDE, ATOVAQUONE
 BEPOTASTINE BESILATE, BEPOTASTINE BESILATE
 BISOPROLOL FUMARATE AND HYDROCHLOROTHIAZIDE, BISOPROLOL FUMARATE
 BREYNA, BUDESONIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** M ****

* MYLAN PHARMACEUTICALS INC
 BROMOCRIPTINE MESYLATE, BROMOCRIPTINE MESYLATE
 BUDESONIDE, BUDESONIDE
 BUSPIRONE HYDROCHLORIDE, BUSPIRONE HYDROCHLORIDE
 CANDESARTAN CILEXETIL AND HYDROCHLOROTHIAZIDE, CANDESARTAN CILEXETIL
 CANDESARTAN CILEXETIL, CANDESARTAN CILEXETIL
 CARBIDOPA AND LEVODOPA, CARBIDOPA
 CARVEDILOL, CARVEDILOL
 CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 CETIRIZINE HYDROCHLORIDE HIVES, CETIRIZINE HYDROCHLORIDE (OTC)
 CHLORTHALIDONE, CHLORTHALIDONE
 CIMETIDINE, CIMETIDINE
 CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE
 CLOZAPINE, CLOZAPINE
 COLCHICINE, COLCHICINE
 CYCLOSPORINE, CYCLOSPORINE
 CYSTAGON, CYSTEAMINE BITARTRATE
 DAPSONE, DAPSONE
 DENAVIR, PENCICLOVIR
 DEXLANSOPRAZOLE, DEXLANSOPRAZOLE
 DIAZEPAM, DIAZEPAM
 DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
 DIMETHYL FUMARATE, DIMETHYL FUMARATE
 DIVALPROEX SODIUM, DIVALPROEX SODIUM
 DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
 ELETRIPTAN HYDROBROMIDE, ELETRIPTAN HYDROBROMIDE
 EMTRICITABINE AND TENOFOVIR DISOPROXIL FUMARATE, EMTRICITABINE
 EPOPROSTENOL SODIUM, EPOPROSTENOL SODIUM
 ERMEZA, LEVOTHYROXINE SODIUM
 ERYGEL, ERYTHROMYCIN
 ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM
 ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM (OTC)
 ESTRADIOL, ESTRADIOL
 ETOPOSIDE, ETOPOSIDE
 EVEROLIMUS, EVEROLIMUS
 FENOFIBRATE, FENOFIBRATE
 FINGOLIMOD HYDROCHLORIDE, FINGOLIMOD HYDROCHLORIDE
 FOSAMPRENAVIR CALCIUM, FOSAMPRENAVIR CALCIUM
 GLATIRAMER ACETATE, GLATIRAMER ACETATE
 HALCINONIDE, HALCINONIDE
 HALOPERIDOL, HALOPERIDOL
 IMATINIB MESYLATE, IMATINIB MESYLATE
 KETOPROFEN, KETOPROFEN
 KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
 LANSOPRAZOLE, LANSOPRAZOLE
 LENALIDOMIDE, LENALIDOMIDE
 LEVETIRACETAM, LEVETIRACETAM
 LEVOTHYROXINE SODIUM, LEVOTHYROXINE SODIUM **
 LISDEXAMFETAMINE DIMESYLATE, LISDEXAMFETAMINE DIMESYLATE
 LOPERAMIDE HYDROCHLORIDE, LOPERAMIDE HYDROCHLORIDE
 LORATADINE, LORATADINE (OTC)
 MECLOFENAMATE SODIUM, MECLOFENAMATE SODIUM
 MERCAPTOPYRINE, MERCAPTOPYRINE
 MESALAMINE, MESALAMINE
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 METHIMAZOLE, METHIMAZOLE
 METHOTREXATE SODIUM, METHOTREXATE SODIUM
 METOLAZONE, METOLAZONE
 METOPROLOL TARTRATE, METOPROLOL TARTRATE
 MIRTAZAPINE, MIRTAZAPINE
 MYCOPHENOLATE MOFETIL, MYCOPHENOLATE MOFETIL
 NEVIRAPINE, NEVIRAPINE
 NISOLDIPINE, NISOLDIPINE
 OLANZAPINE, OLANZAPINE
 PAROXETINE HYDROCHLORIDE, PAROXETINE HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** M ****

- * MYLAN PHARMACEUTICALS INC
 PERPHENAZINE AND AMITRIPTYLINE HYDROCHLORIDE, AMITRIPTYLINE HYDROCHLORIDE
 PHENYTEK, PHENYTOIN SODIUM
 PITAVASTATIN CALCIUM, PITAVASTATIN CALCIUM
 POLYETHYLENE GLYCOL 3350, POLYETHYLENE GLYCOL 3350 (OTC)
 POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
 PRASUGREL, PRASUGREL HYDROCHLORIDE
 PRAZOSIN HYDROCHLORIDE, PRAZOSIN HYDROCHLORIDE
 PREDNISON, PREDNISON
 PROPRANOLOL HYDROCHLORIDE, PROPRANOLOL HYDROCHLORIDE
 REGADENOSON, REGADENOSON
 ROFLUMILAST, ROFLUMILAST
 RUFINAMIDE, RUFINAMIDE
 SAXAGLIPTIN AND METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 SAXAGLIPTIN, SAXAGLIPTIN HYDROCHLORIDE
 SILDENAFIL CITRATE, SILDENAFIL CITRATE
 SORAFENIB TOSYLATE, SORAFENIB TOSYLATE
 SPIRONOLACTONE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 SPIRONOLACTONE, SPIRONOLACTONE
 SUCRALFATE, SUCRALFATE
 SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
 SUNITINIB MALATE, SUNITINIB MALATE
 SYMFI LO, EFAVIRENZ
 TACROLIMUS, TACROLIMUS
 TAMOXIFEN CITRATE, TAMOXIFEN CITRATE
 TELMISARTAN AND AMLODIPINE, AMLODIPINE BESYLATE
 TELMISARTAN, TELMISARTAN
 TETRABENAZINE, TETRABENAZINE
 THIORIDAZINE HYDROCHLORIDE, THIORIDAZINE HYDROCHLORIDE
 TIMOLOL MALEATE, TIMOLOL MALEATE
 TRAVOPROST, TRAVOPROST
 TRETINOIN, TRETINOIN
 TRIFLUOPERAZINE HYDROCHLORIDE, TRIFLUOPERAZINE HYDROCHLORIDE
 VALSARTAN, VALSARTAN
 VARENICLINE TARTRATE, VARENICLINE TARTRATE
 VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE
 VUSION, MICONAZOLE NITRATE
 WIXELA INHUB, FLUTICASONE PROPIONATE
 ZONALON, DOXEPIN HYDROCHLORIDE
- * MYLAN PHARMACEUTICALS INC A VIATRIS CO
 DAPTOMYCIN, DAPTOMYCIN

MYLAN ASI

- * MYLAN ASI LLC
 ACETAZOLAMIDE SODIUM, ACETAZOLAMIDE SODIUM
 ADENOSINE, ADENOSINE
 GRANISETRON HYDROCHLORIDE, GRANISETRON HYDROCHLORIDE

MYLAN INSTITUTIONAL

- * MYLAN INSTITUTIONAL INC
 BUSULFAN, BUSULFAN
- * MYLAN INSTITUTIONAL LLC
 AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE
 ARGATROBAN, ARGATROBAN
 BIVALIRUDIN, BIVALIRUDIN
 CIDOFOVIR, CIDOFOVIR
 DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE
 DIPHENHYDRAMINE HYDROCHLORIDE, DIPHENHYDRAMINE HYDROCHLORIDE
 DURACLON, CLONIDINE HYDROCHLORIDE
 ESMOLOL HYDROCHLORIDE, ESMOLOL HYDROCHLORIDE
 ETHACRYNATE SODIUM, ETHACRYNATE SODIUM
 FOMEPIZOLE, FOMEPIZOLE
 IBUTILIDE FUMARATE, IBUTILIDE FUMARATE
 ISOSULFAN BLUE, ISOSULFAN BLUE
 MELPHALAN HYDROCHLORIDE, MELPHALAN HYDROCHLORIDE
 METHADONE HYDROCHLORIDE, METHADONE HYDROCHLORIDE
 NALOXONE HYDROCHLORIDE, NALOXONE HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** M ******* MYLAN INSTITUTIONAL LLC**

OCTREOTIDE ACETATE (PRESERVATIVE FREE), OCTREOTIDE ACETATE
 PALONOSETRON HYDROCHLORIDE, PALONOSETRON HYDROCHLORIDE
 RIMSO-50, DIMETHYL SULFOXIDE
 ROCURONIUM BROMIDE, ROCURONIUM BROMIDE
 SOTRADECOL, SODIUM TETRADECYL SULFATE
 THIAMINE HYDROCHLORIDE, THIAMINE HYDROCHLORIDE
 TRANEXAMIC ACID, TRANEXAMIC ACID
 ULTIVA, REMIFENTANIL HYDROCHLORIDE

MYLAN IRELAND LTD*** MYLAN IRELAND LTD**

ARIXTRA, FONDAPARINUX SODIUM
 MIACALCIN, CALCITONIN SALMON
 PRETOMANID, PRETOMANID
 YUPELRI, REVEFENACIN

MYLAN LABS LTD*** MYLAN LABORATORIES LTD**

ADENOSINE, ADENOSINE
 BACLOFEN, BACLOFEN
 CIMDUO, LAMIVUDINE
 CLOFARABINE, CLOFARABINE
 CYANOCOBALAMIN, CYANOCOBALAMIN
 DAPTOMYCIN, DAPTOMYCIN
 DEXAMETHASONE SODIUM PHOSPHATE, DEXAMETHASONE SODIUM PHOSPHATE
 DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE
 DOCETAXEL, DOCETAXEL
 DOXORUBICIN HYDROCHLORIDE, DOXORUBICIN HYDROCHLORIDE
 DOXYCYCLINE, DOXYCYCLINE HYCLATE
 EPTIFIBATIDE, EPTIFIBATIDE
 ESMOLOL HYDROCHLORIDE, ESMOLOL HYDROCHLORIDE
 ETOMIDATE, ETOMIDATE
 FAMOTIDINE PRESERVATIVE FREE, FAMOTIDINE
 FAMOTIDINE, FAMOTIDINE
 FLUPHENAZINE DECANOATE, FLUPHENAZINE DECANOATE
 FOSAPREPITANT DIMEGLUMINE, FOSAPREPITANT DIMEGLUMINE
 GEMCITABINE HYDROCHLORIDE, GEMCITABINE HYDROCHLORIDE
 GLUCAGON, GLUCAGON
 HALOPERIDOL DECANOATE, HALOPERIDOL DECANOATE
 HALOPERIDOL, HALOPERIDOL LACTATE
 HEPARIN SODIUM, HEPARIN SODIUM
 IBANDRONATE SODIUM, IBANDRONATE SODIUM
 LEUCOVORIN CALCIUM PRESERVATIVE FREE, LEUCOVORIN CALCIUM
 LEVETIRACETAM, LEVETIRACETAM
 LINEZOLID, LINEZOLID
 LOPINAVIR AND RITONAVIR, LOPINAVIR
 MAGNESIUM SULFATE IN DEXTROSE 5% IN PLASTIC CONTAINER, MAGNESIUM SULFATE
 MAGNESIUM SULFATE IN PLASTIC CONTAINER, MAGNESIUM SULFATE
 MOXIFLOXACIN HYDROCHLORIDE IN SODIUM CHLORIDE 0.8% IN PLASTIC CONTAINER, MOXIFLOXACIN
 MYCOPHENOLATE MOFETIL HYDROCHLORIDE, MYCOPHENOLATE MOFETIL HYDROCHLORIDE
 NALOXONE HYDROCHLORIDE, NALOXONE HYDROCHLORIDE
 NOREPINEPHRINE BITARTRATE, NOREPINEPHRINE BITARTRATE
 OXALIPLATIN, OXALIPLATIN
 PAMIDRONATE DISODIUM, PAMIDRONATE DISODIUM
 POSACONAZOLE, POSACONAZOLE
 PROCHLORPERAZINE EDISYLATE, PROCHLORPERAZINE EDISYLATE
 RIFAMPIN, RIFAMPIN
 ROPIVACAINE HYDROCHLORIDE, ROPIVACAINE HYDROCHLORIDE
 SODIUM NITROPRUSSIDE, SODIUM NITROPRUSSIDE
 SULFAMETHOXAZOLE AND TRIMETHOPRIM, SULFAMETHOXAZOLE
 SYMFI, EFAVIRENZ
 TOBRAMYCIN SULFATE, TOBRAMYCIN SULFATE
 TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE
 VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE
 VECURONIUM BROMIDE, VECURONIUM BROMIDE
 ZOLEDRONIC ACID, ZOLEDRONIC ACID

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** M ******MYLAN PHARMS INC**

- * MYLAN PHARMACEUTICALS INC
 - ABACAVIR SULFATE, ABACAVIR SULFATE
 - ACYCLOVIR, ACYCLOVIR
 - AMNESTEEM, ISOTRETINOIN
 - ARMODAFINIL, ARMODAFINIL
 - ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM
 - AVITA, TRETINOIN
 - CHLORDIAZEPOXIDE AND AMITRIPTYLINE HYDROCHLORIDE, AMITRIPTYLINE HYDROCHLORIDE
 - CLINDAMYCIN PHOSPHATE AND BENZOYL PEROXIDE, BENZOYL PEROXIDE
 - DOXEPIN HYDROCHLORIDE, DOXEPIN HYDROCHLORIDE
 - DOXYLAMINE SUCCINATE AND PYRIDOXINE HYDROCHLORIDE, DOXYLAMINE SUCCINATE
 - ESZOPICLONE, ESZOPICLONE
 - FENOFIBRATE, FENOFIBRATE
 - LANSOPRAZOLE, LANSOPRAZOLE
 - LITHIUM CARBONATE, LITHIUM CARBONATE
 - METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 - METOPROLOL SUCCINATE, METOPROLOL SUCCINATE
 - MIDODRINE HYDROCHLORIDE, MIDODRINE HYDROCHLORIDE
 - NEVIRAPINE, NEVIRAPINE
 - PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
 - RILUZOLE, RILUZOLE
 - RIZATRIPTAN BENZOATE, RIZATRIPTAN BENZOATE
 - VALACYCLOVIR HYDROCHLORIDE, VALACYCLOVIR HYDROCHLORIDE
 - VALSARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 - VORICONAZOLE, VORICONAZOLE
- * MYLAN PHARMACEUTICALS INC.
 - FLUVASTATIN SODIUM, FLUVASTATIN SODIUM

MYLAN SPCLT VIATRIS

- * MYLAN SPECIALTY LP A VIATRIS CO
 - DIPENTUM, OLSALAZINE SODIUM

MYLAN SPECIALITY LP

- * MYLAN SPECIALTY LP
 - CORTIFOAM, HYDROCORTISONE ACETATE
 - DEPEN, PENICILLAMINE
 - DYMISTA, AZELASTINE HYDROCHLORIDE
 - EDLUAR, ZOLPIDEM TARTRATE
 - ELESTRIN, ESTRADIOL
 - EPIFOAM, HYDROCORTISONE ACETATE
 - EPIPEN JR., EPINEPHRINE
 - EPIPEN, EPINEPHRINE
 - FELBATOL, FELBAMATE
 - GASTROCROM, CROMOLYN SODIUM
 - LEVALBUTEROL HYDROCHLORIDE, LEVALBUTEROL HYDROCHLORIDE
 - PROCTOFOAM HC, HYDROCORTISONE ACETATE
 - ROWASA, MESALAMINE
 - SFROWASA, MESALAMINE
 - SOMA, CARISOPRODOL
 - TOBI PODHALER, TOBRAMYCIN
 - TOBI, TOBRAMYCIN

MYLAN SPECLT

- * MYLAN SPECIALTY LP
 - PERFORMIST, FORMOTEROL FUMARATE

MYLAN TECH VIATRIS

- * MYLAN TECHNOLOGIES INC A VIATRIS CO
 - METHYLPHENIDATE, METHYLPHENIDATE

MYLAN TECHNOLOGIES

- * MYLAN TECHNOLOGIES INC
 - BUPRENORPHINE HYDROCHLORIDE AND NALOXONE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE
 - CLONIDINE, CLONIDINE
 - ESTRADIOL, ESTRADIOL
 - FENTANYL-100, FENTANYL
 - FENTANYL-12, FENTANYL
 - FENTANYL-25, FENTANYL
 - FENTANYL-37, FENTANYL

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** M ****

* MYLAN TECHNOLOGIES INC
 FENTANYL-50, FENTANYL
 FENTANYL-62, FENTANYL
 FENTANYL-75, FENTANYL
 FENTANYL-87, FENTANYL
 LIDOCAINE, LIDOCAINE
 NITROGLYCERIN, NITROGLYCERIN
 RIVASTIGMINE, RIVASTIGMINE
 SCOPOLAMINE, SCOPOLAMINE
 XULANE, ETHINYL ESTRADIOL

**** N ******NAARI PTE LTD**

* NAARI PTE LTD
 DESOGESTREL AND ETHINYL ESTRADIOL, DESOGESTREL
 DROSPIRENONE AND ETHINYL ESTRADIOL, DROSPIRENONE
 ESTRADIOL AND NORETHINDRONE ACETATE, ESTRADIOL
 LEVONORGESTREL AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
 LEVONORGESTREL, LEVONORGESTREL (OTC)
 NORETHINDRONE AND ETHINYL ESTRADIOL AND FERROUS FUMARATE, ETHINYL ESTRADIOL
 NORETHINDRONE AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
 NORETHINDRONE, NORETHINDRONE
 NORGESTIMATE AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL

NAL PHARM

* NAL PHARMACEUTICAL GROUP LTD
 LIDOCAINE, LIDOCAINE

NALPROPION

* NALPROPION PHARMACEUTICALS LLC
 CONTRAVE, BUPROPION HYDROCHLORIDE

NANG KUANG PHARM CO

* NANG KUANG PHARMACEUTICAL CO LTD
 ICATIBANT ACETATE, ICATIBANT ACETATE
 LINEZOLID, LINEZOLID
 PEMETREXED DISODIUM, PEMETREXED DISODIUM

NANJING

* NANJING SIMCERE DONGYUAN PHARMACEUTICAL CO LTD
 CELECOXIB, CELECOXIB

NANJING DAOQUN

* NANJING DAOQUN PHARMACEUTICAL R AND D CO LTD
 OSELTAMIVIR PHOSPHATE, OSELTAMIVIR PHOSPHATE

NANJING KING-FRIEND

* NANJING KING-FRIEND BIOCHEMICAL PHARMACEUTICAL CO LTD
 CYANOCOBALAMIN, CYANOCOBALAMIN
 ENOXAPARIN SODIUM (PRESERVATIVE FREE), ENOXAPARIN SODIUM
 ENOXAPARIN SODIUM, ENOXAPARIN SODIUM
 HEPARIN SODIUM PRESERVATIVE FREE, HEPARIN SODIUM
 HEPARIN SODIUM, HEPARIN SODIUM

NANOCOPOEIA

* NANOCOPOEIA LLC
 PHYRAGO, DASATINIB

NAPO PHARMS INC

* NAPO PHARMACEUTICALS INC
 MYTESI, CROFELEMER

NATCO

* NATCO PHARMA LTD
 ALPRAZOLAM, ALPRAZOLAM
 CARISOPRODOL, CARISOPRODOL
 GEFITINIB, GEFITINIB
 GLYCOPYRROLATE, GLYCOPYRROLATE
 LANSOPRAZOLE, LANSOPRAZOLE (OTC)
 NITROGLYCERIN, NITROGLYCERIN
 OSELTAMIVIR PHOSPHATE, OSELTAMIVIR PHOSPHATE
 TERIFLUNOMIDE, TERIFLUNOMIDE
 TIPIRACIL HYDROCHLORIDE AND TRIFLURIDINE, TIPIRACIL HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** N ******NATCO PHARMA**

- * NATCO PHARMA LTD
GRANISETRON HYDROCHLORIDE, GRANISETRON HYDROCHLORIDE
SUNITINIB MALATE, SUNITINIB MALATE

NATCO PHARMA LTD

- * NATCO PHARMA LIMITED
CHLOROQUINE PHOSPHATE, CHLOROQUINE PHOSPHATE
- * NATCO PHARMA LTD
ANASTROZOLE, ANASTROZOLE
ARMODAFINIL, ARMODAFINIL
AZACITIDINE, AZACITIDINE
CHLOROQUINE PHOSPHATE, CHLOROQUINE PHOSPHATE
ERLOTINIB HYDROCHLORIDE, ERLOTINIB HYDROCHLORIDE
IMATINIB MESYLATE, IMATINIB MESYLATE
LANSOPRAZOLE, LANSOPRAZOLE
LANTHANUM CARBONATE, LANTHANUM CARBONATE
LAPATINIB DITOSYLATE, LAPATINIB DITOSYLATE
LETROZOLE, LETROZOLE
ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
RIZATRIPTAN BENZOATE, RIZATRIPTAN BENZOATE
TRIHEXYPHENIDYL HYDROCHLORIDE, TRIHEXYPHENIDYL HYDROCHLORIDE

NAVINTA LLC

- * NAVINTA LLC
BENZTROPINE MESYLATE, BENZTROPINE MESYLATE
CARGLUMIC ACID, CARGLUMIC ACID
CARMUSTINE, CARMUSTINE
FAMOTIDINE, FAMOTIDINE
FOMEPIZOLE, FOMEPIZOLE
HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
PENICILLAMINE, PENICILLAMINE
RIBAVIRIN, RIBAVIRIN
ROPIVACAINE HYDROCHLORIDE, ROPIVACAINE HYDROCHLORIDE
SODIUM PHENYLACETATE AND SODIUM BENZOATE, SODIUM BENZOATE
TRIENTINE HYDROCHLORIDE, TRIENTINE HYDROCHLORIDE

NCM USA BRONX LLC

- * NCM USA BRONX LLC
AMMONIA N 13, AMMONIA N-13
FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18
SODIUM FLUORIDE F-18, SODIUM FLUORIDE F-18

NE RX PHARMA

- * NE RX PHARMA LLC
ACETAZOLAMIDE, ACETAZOLAMIDE
BUTALBITAL AND ACETAMINOPHEN, ACETAMINOPHEN
METHSCOPOLAMINE BROMIDE, METHSCOPOLAMINE BROMIDE
METOLAZONE, METOLAZONE
VARENICLINE TARTRATE, VARENICLINE TARTRATE

NEOS THERAPS

- * NEOS THERAPEUTICS
ADZENYS XR-ODT, AMPHETAMINE

NEOS THERAPS INC

- * NEOS THERAPEUTICS INC
COTEMPLA XR-ODT, METHYLPHENIDATE

NEPHRON

- * NEPHRON CORP
ALBUTEROL SULFATE, ALBUTEROL SULFATE
IPRATROPIUM BROMIDE, IPRATROPIUM BROMIDE
- * NEPHRON PHARMACEUTICALS CORP
ALBUTEROL SULFATE AND IPRATROPIUM BROMIDE, ALBUTEROL SULFATE
ALBUTEROL SULFATE, ALBUTEROL SULFATE
- * NEPHRON PHARMACEUTICALS CORPORATION
BUDESONIDE, BUDESONIDE
KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, SODIUM CHLORIDE
STERILE WATER FOR INJECTION, STERILE WATER FOR INJECTION
VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** N ******NEURELIS INC**

* NEURELIS INC
VALTOCO, DIAZEPAM

NEUROCRINE

* NEUROCRINE BIOSCIENCES INC
CRENESSITY, CRINECERFONT
INGREZZA SPRINKLE, VALBENAZINE TOSYLATE
INGREZZA, VALBENAZINE TOSYLATE

NEW HEIGHTSRX

* NEW HEIGHTSRX LLC
CLEMASTINE FUMARATE, CLEMASTINE FUMARATE

NEXTSOURCE

* NEXTSOURCE BIOTECHNOLOGY LLC
CYCLOPHOSPHAMIDE, CYCLOPHOSPHAMIDE

NEXTWAVE

* NEXTWAVE PHARMACEUTICALS INC A SUB OF TRIS PHARMA INC
QUILLIVANT XR, METHYLPHENIDATE HYDROCHLORIDE

NEXTWAVE PHARMS

* NEXTWAVE PHARMACEUTICALS INC
QUILLICHEW ER, METHYLPHENIDATE HYDROCHLORIDE

NEXUS

* NEXUS PHARMACEUTICALS LLC
ARSENIC TRIOXIDE, ARSENIC TRIOXIDE
COLISTIMETHATE SODIUM, COLISTIMETHATE SODIUM
CYCLOPHOSPHAMIDE, CYCLOPHOSPHAMIDE
DICYCLOMINE HYDROCHLORIDE, DICYCLOMINE HYDROCHLORIDE
EMERPHED, EPHEDRINE SULFATE
ERYTHROMYCIN LACTOBIONATE, ERYTHROMYCIN LACTOBIONATE
ISOPROTERENOL HYDROCHLORIDE, ISOPROTERENOL HYDROCHLORIDE
LEVETIRACETAM IN SODIUM CHLORIDE, LEVETIRACETAM
METHYLENE BLUE, METHYLENE BLUE
NELARABINE, NELARABINE
POTASSIUM CHLORIDE 10MEQ, POTASSIUM CHLORIDE
POTASSIUM CHLORIDE 20MEQ, POTASSIUM CHLORIDE
POTASSIUM CHLORIDE 40MEQ, POTASSIUM CHLORIDE
PROCAINAMIDE HYDROCHLORIDE, PROCAINAMIDE HYDROCHLORIDE
PROCHLORPERAZINE EDISYLATE, PROCHLORPERAZINE EDISYLATE
SODIUM NITROPRUSSIDE, SODIUM NITROPRUSSIDE
STERILE WATER FOR INJECTION, STERILE WATER FOR INJECTION
TESTOSTERONE ENANTHATE, TESTOSTERONE ENANTHATE
TIROFIBAN HYDROCHLORIDE, TIROFIBAN HYDROCHLORIDE
TRANEXAMIC ACID, TRANEXAMIC ACID

NEXUS PHARMS

* NEXUS PHARMACEUTICALS INC
FLUORESCEIN SODIUM, FLUORESCEIN SODIUM
MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE
POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
SODIUM CHLORIDE 0.9%, SODIUM CHLORIDE

NIAGARA PHARMS

* NIAGARA PHARMACEUTICALS INC
PUR-WASH, PURIFIED WATER (OTC)

NIPPON SHINYAKU

* NIPPON SHINYAKU CO LTD
VILTEPSO, VILTOLARSEN

NIVAGEN PHARMS INC

* NIVAGEN PHARMACEUTICALS INC
CALCIUM GLUCONATE, CALCIUM GLUCONATE
DECITABINE, DECITABINE
REMIFENTANIL HYDROCHLORIDE, REMIFENTANIL HYDROCHLORIDE
ZINC SULFATE, ZINC SULFATE

NOBELPHARMA

* NOBELPHARMA CO LTD
HYFTOR, SIROLIMUS

NODEN PHARMA

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** N ******* NODEN PHARMA DAC**

GOPRELTO, COCAINE HYDROCHLORIDE
TEKTURNA, ALISKIREN HEMIFUMARATE

NORDIC PHARMA*** NORDIC PHARMA INC**

NEOMYCIN AND POLYMYXIN B SULFATES AND GRAMICIDIN, GRAMICIDIN

NORTEC DEV ASSOC*** NORTEC DEVELOPMENT ASSOC INC**

PROPRANOLOL HYDROCHLORIDE, PROPRANOLOL HYDROCHLORIDE

NORTHLAND*** NORTHLAND NUCLEAR MEDICINE LLC**

FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18

NORTHSTAR HLTHCARE*** NORTHSTAR HEALTHCARE HOLDINGS LTD**

ALLOPURINOL, ALLOPURINOL
BACLOFEN, BACLOFEN
GEMFIBROZIL, GEMFIBROZIL
HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE
PROPRANOLOL HYDROCHLORIDE, PROPRANOLOL HYDROCHLORIDE

NORTON WATERFORD*** NORTON WATERFORD LTD**

QVAR REDIHALER, BECLOMETHASONE DIPROPIONATE

NORVIUM BIOSCIENCE*** NORVIUM BIOSCIENCE LLC**

DEFERASIROX, DEFERASIROX
RITONAVIR, RITONAVIR
SODIUM NITROPRUSSIDE, SODIUM NITROPRUSSIDE
TENOFVIR DISOPROXIL FUMARATE, TENOFVIR DISOPROXIL FUMARATE

NORWICH*** NORWICH PHARMACEUTICALS INC**

LISDEXAMFETAMINE DIMESYLATE, LISDEXAMFETAMINE DIMESYLATE

NOSTRUM LABS INC*** NOSTRUM LABORATORIES INC**

ACETAZOLAMIDE, ACETAZOLAMIDE
BUTALBITAL, ASPIRIN AND CAFFEINE, ASPIRIN
CALCIUM ACETATE, CALCIUM ACETATE
CARBAMAZEPINE, CARBAMAZEPINE
CARISOPRODOL, CARISOPRODOL
CLARITHROMYCIN, CLARITHROMYCIN
DAPSONE, DAPSONE
ELIXOPHYLLIN, THEOPHYLLINE
FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
NITROFURANTOIN, NITROFURANTOIN
OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN
PINDOLOL, PINDOLOL
PIROXICAM, PIROXICAM
PROMETHAZINE HYDROCHLORIDE AND CODEINE PHOSPHATE, CODEINE PHOSPHATE
PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
SUCRALFATE, SUCRALFATE
THEOPHYLLINE, THEOPHYLLINE
VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE

NOSTRUM PHARMS LLC*** NOSTRUM PHARMACEUTICALS LLC**

METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
THEOPHYLLINE, THEOPHYLLINE

NOVA LABS LTD*** NOVA LABORATORIES LTD**

PURIXAN, MERCAPTOPYRINE
XROMI, HYDROXYUREA

NOVADAQ TECH*** NOVADAQ TECHNOLOGIES ULC**

SPY AGENT GREEN KIT, INDOCYANINE GREEN

NOVARTIS

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** N ****

* NOVARTIS PHARMACEUTICALS CORP
 AFINITOR, EVEROLIMUS
 ALOMIDE, LODOXAMIDE TROMETHAMINE
 BETOPTIC S, BETAXOLOL HYDROCHLORIDE
 COARTEM, ARTEMETHER
 DESFERAL, DEFEROXAMINE MESYLATE
 DIOVAN HCT, HYDROCHLOROTHIAZIDE
 DIOVAN, VALSARTAN
 EGATEN, TRICLABENDAZOLE
 ENTRESTO SPRINKLE, SACUBITRIL
 EXFORGE HCT, AMLODIPINE BESYLATE
 EXFORGE, AMLODIPINE BESYLATE
 EXJADE, DEFERASIROX
 FABHALTA, IPTACOPAN HYDROCHLORIDE
 GILENYA, FINGOLIMOD HYDROCHLORIDE
 GLEEVEC, IMATINIB MESYLATE
 JADENU SPRINKLE, DEFERASIROX
 KISQALI FEMARA CO-PACK (COPACKAGED), LETROZOLE
 KISQALI, RIBOCICLIB SUCCINATE
 LEQVIO, INCLISIRAN SODIUM
 LOCAMETZ, GALLIUM GA-68 GOZETOTIDE
 MAYZENT, SIPONIMOD
 MEKINIST, TRAMETINIB DIMETHYL SULFOXIDE
 MYFORTIC, MYCOPHENOLIC SODIUM
 NEORAL, CYCLOSPORINE
 PIQRAY, ALPELISIB
 PLUVICTO, LUTETIUM LU-177 VIPIVOTIDE TETRAKETAN
 PROMACTA KIT, ELTROMBOPAG OLAMINE
 PROMACTA, ELTROMBOPAG OLAMINE
 RYDAPT, MIDOSTAURIN
 SANDIMMUNE, CYCLOSPORINE
 SANDOSTATIN LAR, OCTREOTIDE ACETATE
 SANDOSTATIN, OCTREOTIDE ACETATE
 SCEMBLIX, ASCIMINIB HYDROCHLORIDE
 TAFINLAR, DABRAFENIB MESYLATE
 TASIGNA, NILOTINIB HYDROCHLORIDE
 TEGRETOL, CARBAMAZEPINE
 TEGRETOL-XR, CARBAMAZEPINE
 TOBRADEX, DEXAMETHASONE
 TOBREX, TOBRAMYCIN
 TRILEPTAL, OXCARBAZEPINE
 TYKERB, LAPATINIB DITOSYLATE
 VIJOICE, ALPELISIB
 VOTRIENT, PAZOPANIB HYDROCHLORIDE
 ZORTRESS, EVEROLIMUS
 ZYKADIA, CERITINIB

NOVARTIS PHARM

* NOVARTIS PHARMACEUTICAL CORP
 AFINITOR DISPERZ, EVEROLIMUS
 TABRECTA, CAPMATINIB HYDROCHLORIDE

NOVARTIS PHARMS

* NOVARTIS PHARMACEUTICALS CORP
 FEMARA, LETROZOLE

NOVARTIS PHARMS CORP

* NOVARTIS PHARMACEUTICALS CORP
 ENTRESTO, SACUBITRIL
 JADENU, DEFERASIROX

NOVAST LABS

* NOVAST LABORATORIES CHINA LTD
 NORETHINDRONE, NORETHINDRONE
 * NOVAST LABORATORIES INC
 DOCETAXEL, DOCETAXEL
 MESALAMINE, MESALAMINE
 * NOVAST LABORATORIES LTD
 ACETAZOLAMIDE, ACETAZOLAMIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** N ******* NOVAST LABORATORIES LTD**

BISMUTH SUBCITRATE POTASSIUM, METRONIDAZOLE AND TETRACYCLINE HYDROCHLORIDE, BISMUTH
 BUTALBITAL, ACETAMINOPHEN AND CAFFEINE, ACETAMINOPHEN
 CARBAMAZEPINE, CARBAMAZEPINE
 CARBOPLATIN, CARBOPLATIN
 CARISOPRODOL, CARISOPRODOL
 CHABELINA FE, ETHINYL ESTRADIOL
 CHLORTHALIDONE, CHLORTHALIDONE
 CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
 CYCLOBENZAPRINE HYDROCHLORIDE, CYCLOBENZAPRINE HYDROCHLORIDE
 CYPROHEPTADINE HYDROCHLORIDE, CYPROHEPTADINE HYDROCHLORIDE
 DECITABINE, DECITABINE
 DESIPRAMINE HYDROCHLORIDE, DESIPRAMINE HYDROCHLORIDE
 DESMOPRESSIN ACETATE, DESMOPRESSIN ACETATE
 DESOGESTREL AND ETHINYL ESTRADIOL, DESOGESTREL
 DICLOFENAC POTASSIUM, DICLOFENAC POTASSIUM
 DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
 DOLISHALE, ETHINYL ESTRADIOL
 ESTRADIOL AND NORETHINDRONE ACETATE, ESTRADIOL
 GEMCITABINE HYDROCHLORIDE, GEMCITABINE HYDROCHLORIDE
 HER STYLE, LEVONORGESTREL (OTC)
 INDOMETHACIN, INDOMETHACIN
 IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE
 LARIN 1.5/30, ETHINYL ESTRADIOL
 LARIN 1/20, ETHINYL ESTRADIOL
 LARIN 24 FE, ETHINYL ESTRADIOL
 LARIN FE 1.5/30, ETHINYL ESTRADIOL
 LARIN FE 1/20, ETHINYL ESTRADIOL
 LERIBANE, ETHINYL ESTRADIOL
 LEUCOVORIN CALCIUM, LEUCOVORIN CALCIUM
 LO-MALMOREDE, ETHINYL ESTRADIOL
 MALMOREDE, ETHINYL ESTRADIOL
 MELAMISA, DROSPIRENONE
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 METHENAMINE HIPPURATE, METHENAMINE HIPPURATE
 METOPROLOL SUCCINATE, METOPROLOL SUCCINATE
 MEXILETINE HYDROCHLORIDE, MEXILETINE HYDROCHLORIDE
 NIFEDIPINE, NIFEDIPINE
 NORETHINDRONE ACETATE, NORETHINDRONE ACETATE
 NORETHINDRONE, NORETHINDRONE
 OXALIPLATIN, OXALIPLATIN
 OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE
 PIMTREA, DESOGESTREL
 PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE
 PRIMAQUINE PHOSPHATE, PRIMAQUINE PHOSPHATE
 PROBENECID AND COLCHICINE, COLCHICINE
 QUETIAPINE FUMARATE, QUETIAPINE FUMARATE
 QUININE SULFATE, QUININE SULFATE
 RANOLAZINE, RANOLAZINE
 SETLAKIN, ETHINYL ESTRADIOL
 TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE
 TOLCAPONE, TOLCAPONE
 TOPOTECAN HYDROCHLORIDE, TOPOTECAN HYDROCHLORIDE
 TRI-LO-LINYAH, ETHINYL ESTRADIOL
 TRIAZOLAM, TRIAZOLAM
 YAELA, DROSPIRENONE

NOVAST LABS LTD*** NOVAST LABORATORIES LTD**

DASETTA 1/35, ETHINYL ESTRADIOL
 DASETTA 7/7/7, ETHINYL ESTRADIOL
 ELINEST, ETHINYL ESTRADIOL
 FALMINA, ETHINYL ESTRADIOL
 LEVONEST, ETHINYL ESTRADIOL
 MONO-LINYAH, ETHINYL ESTRADIOL
 PHILITH, ETHINYL ESTRADIOL

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** N ****

* NOVAST LABORATORIES LTD
 TRI-LINYAH, ETHINYL ESTRADIOL
 WERA, ETHINYL ESTRADIOL

NOVATECH SA

* NOVATECH SA
 STERITALC, TALC

NOVEL LABS INC

* NOVEL LABORATORIES INC
 AMLODIPINE BESYLATE AND VALSARTAN, AMLODIPINE BESYLATE
 CARBIDOPA, CARBIDOPA
 CEVIMELINE HYDROCHLORIDE, CEVIMELINE HYDROCHLORIDE
 CHILDREN'S CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
 DESOXIMETASONE, DESOXIMETASONE
 DEXMETHYLPHENIDATE HYDROCHLORIDE, DEXMETHYLPHENIDATE HYDROCHLORIDE
 DEXTROAMPHETAMINE SULFATE, DEXTROAMPHETAMINE SULFATE
 DIAZEPAM, DIAZEPAM
 DICLOFENAC SODIUM, DICLOFENAC SODIUM
 DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
 FAMOTIDINE, FAMOTIDINE
 FLUCYTOSINE, FLUCYTOSINE
 FLUOCINOLONE ACETONIDE, FLUOCINOLONE ACETONIDE
 FLUOCINONIDE, FLUOCINONIDE
 HYDROCODONE BITARTRATE AND HOMATROPINE METHYLBROMIDE, HOMATROPINE METHYLBROMIDE
 LEVONORGESTREL, LEVONORGESTREL (OTC)
 LINEZOLID, LINEZOLID
 METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
 METOCLOPRAMIDE HYDROCHLORIDE, METOCLOPRAMIDE HYDROCHLORIDE
 MONOBASIC SODIUM PHOSPHATE AND DIBASIC SODIUM PHOSPHATE, SODIUM PHOSPHATE, DIBASIC,
 MORPHINE SULFATE, MORPHINE SULFATE
 NITROFURANTOIN, NITROFURANTOIN
 NITROFURANTOIN, NITROFURANTOIN, MACROCRYSTALLINE
 OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN
 OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
 PEG 3350 AND ELECTROLYTES, POLYETHYLENE GLYCOL 3350
 PEG-3350, POTASSIUM CHLORIDE, SODIUM BICARBONATE, SODIUM CHLORIDE, POLYETHYLENE GLYCOL
 PEG-3350, SODIUM SULFATE, SODIUM CHLORIDE, POTASSIUM CHLORIDE, SODIUM ASCORBATE AND
 PHENELZINE SULFATE, PHENELZINE SULFATE
 POLYETHYLENE GLYCOL 3350, POLYETHYLENE GLYCOL 3350 (OTC)
 POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
 SODIUM SULFATE, POTASSIUM SULFATE AND MAGNESIUM SULFATE, MAGNESIUM SULFATE
 TEMAZEPAM, TEMAZEPAM
 TRIMETHOPRIM, TRIMETHOPRIM
 VORICONAZOLE, VORICONAZOLE
 ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE

NOVELGENIX THERAPS

* NOVELGENIX THERAPEUTICS PVT LTD
 NAPROXEN SODIUM, NAPROXEN SODIUM (OTC)

NOVEN

* NOVEN PHARMACEUTICALS INC
 MINIVELLE, ESTRADIOL

NOVEN PHARMS INC

* NOVEN PHARMACEUTICALS INC
 COMBIPATCH, ESTRADIOL
 DAYTRANA, METHYLPHENIDATE
 XELSTRYM, DEXTROAMPHETAMINE

NOVITIUM PHARMA

* NOVITIUM PHARMA LLC
 ACETAZOLAMIDE, ACETAZOLAMIDE
 ACYCLOVIR, ACYCLOVIR
 ALENDRONATE SODIUM, ALENDRONATE SODIUM
 ALPRAZOLAM, ALPRAZOLAM
 ATENOLOL AND CHLORTHALIDONE, ATENOLOL
 BETAINE, BETAINE
 BISOPROLOL FUMARATE AND HYDROCHLOROTHIAZIDE, BISOPROLOL FUMARATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** N ****

* NOVITIUM PHARMA LLC
 BISOPROLOL FUMARATE, BISOPROLOL FUMARATE
 CARBAMAZEPINE, CARBAMAZEPINE
 CARGLUMIC ACID, CARGLUMIC ACID
 CHLORZOXAZONE, CHLORZOXAZONE
 CLORAZEPATE DIPOTASSIUM, CLORAZEPATE DIPOTASSIUM
 CLOTRIMAZOLE, CLOTRIMAZOLE
 DAPSONE, DAPSONE
 DEXAMETHASONE, DEXAMETHASONE
 DIAZOXIDE, DIAZOXIDE
 DICYCLOMINE HYDROCHLORIDE, DICYCLOMINE HYDROCHLORIDE
 DIGOXIN, DIGOXIN
 ESTAZOLAM, ESTAZOLAM
 ESTRADIOL, ESTRADIOL
 FAMOTIDINE, FAMOTIDINE
 FELBAMATE, FELBAMATE
 FLUDROCORTISONE ACETATE, FLUDROCORTISONE ACETATE
 FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 FLUPHENAZINE HYDROCHLORIDE, FLUPHENAZINE HYDROCHLORIDE
 INDOMETHACIN, INDOMETHACIN
 KETOCONAZOLE, KETOCONAZOLE
 L-GLUTAMINE, L-GLUTAMINE
 LACOSAMIDE, LACOSAMIDE
 LEVOCARNITINE SF, LEVOCARNITINE
 LEVOCARNITINE, LEVOCARNITINE
 LEVOFLOXACIN, LEVOFLOXACIN
 LEVORPHANOL TARTRATE, LEVORPHANOL TARTRATE
 LIDOCAINE HYDROCHLORIDE, LIDOCAINE HYDROCHLORIDE
 MEGESTROL ACETATE, MEGESTROL ACETATE
 MELOXICAM, MELOXICAM
 METHSUXIMIDE, METHSUXIMIDE
 METHYLTESTOSTERONE, METHYLTESTOSTERONE
 NABUMETONE, NABUMETONE
 NAPROXEN, NAPROXEN
 NITISINONE, NITISINONE
 NITROFURANTOIN, NITROFURANTOIN
 OXYBUTYNYN CHLORIDE, OXYBUTYNYN CHLORIDE
 PAROXETINE HYDROCHLORIDE, PAROXETINE HYDROCHLORIDE
 PHENOXYBENZAMINE HYDROCHLORIDE, PHENOXYBENZAMINE HYDROCHLORIDE
 POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
 PRAZOSIN HYDROCHLORIDE, PRAZOSIN HYDROCHLORIDE
 PREDNISONE, PREDNISONE
 PROCHLORPERAZINE MALEATE, PROCHLORPERAZINE MALEATE
 PRUCALOPRIDE SUCCINATE, PRUCALOPRIDE SUCCINATE
 PYRAZINAMIDE, PYRAZINAMIDE
 PYRIDOSTIGMINE BROMIDE, PYRIDOSTIGMINE BROMIDE
 RIFABUTIN, RIFABUTIN
 SELEGILINE HYDROCHLORIDE, SELEGILINE HYDROCHLORIDE
 SILDENAFIL CITRATE, SILDENAFIL CITRATE
 SIROLIMUS, SIROLIMUS
 SOVUNA, HYDROXYCHLOROQUINE SULFATE
 SULFAMETHOXAZOLE AND TRIMETHOPRIM, SULFAMETHOXAZOLE
 TADALAFIL, TADALAFIL
 TEZRULY, TERAZOSIN HYDROCHLORIDE
 THIOTHIXENE, THIOTHIXENE
 TRANLYCYPROMINE SULFATE, TRANLYCYPROMINE SULFATE
 TRIHEXYPHENIDYL HYDROCHLORIDE, TRIHEXYPHENIDYL HYDROCHLORIDE
 TRIMETHOPRIM, TRIMETHOPRIM
 VALSARTAN, VALSARTAN

NOVO

* NOVO NORDISK INC
 OZEMPIC, SEMAGLUTIDE
 RIVFLOZA, NEDOSIRAN SODIUM
 RYBELSUS, SEMAGLUTIDE
 SAXENDA, LIRAGLUTIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** N ****

* NOVO NORDISK INC
WEGOVY, SEMAGLUTIDE

NOVO NORDISK INC

* NOVO NORDISK INC
VAGIFEM, ESTRADIOL
VICTOZA, LIRAGLUTIDE

NOVUGEN

* NOVUGEN ONCOLOGY SDN BHD
ABIRATERONE ACETATE, ABIRATERONE ACETATE
LENALIDOMIDE, LENALIDOMIDE
PAZOPANIB HYDROCHLORIDE, PAZOPANIB HYDROCHLORIDE
* NOVUGEN PHARMA MALAYSIA SDN BHD
MIDODRINE HYDROCHLORIDE, MIDODRINE HYDROCHLORIDE

NUKEMED

* NUKEMED INC DBA SPECTRONRX
AMMONIA N 13, AMMONIA N-13
FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18
SODIUM FLUORIDE F-18, SODIUM FLUORIDE F-18

NUVO PHARM

* NUVO PHARMACEUTICAL INC
BENZTROPINE MESYLATE, BENZTROPINE MESYLATE
DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE
DEXTROAMPHETAMINE SULFATE, DEXTROAMPHETAMINE SULFATE
DIAZEPAM, DIAZEPAM
OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
PHENTERMINE HYDROCHLORIDE, PHENTERMINE HYDROCHLORIDE

NUVO PHARMS INC

* NUVO PHAMACEUTICALS INC
BUTALBITAL, ACETAMINOPHEN AND CAFFEINE, ACETAMINOPHEN
CHLORDIAZEPOXIDE HYDROCHLORIDE AND CLIDINIUM BROMIDE, CHLORDIAZEPOXIDE HYDROCHLORIDE
FOLIC ACID, FOLIC ACID
HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE
NAPROXEN, NAPROXEN
POLYETHYLENE GLYCOL 3350, POLYETHYLENE GLYCOL 3350 (OTC)
SODIUM POLYSTYRENE SULFONATE, SODIUM POLYSTYRENE SULFONATE
SULFASALAZINE, SULFASALAZINE

NXDC

* NX DEVELOPMENT CORP
GLEOLAN, AMINOLEVULINIC ACID HYDROCHLORIDE

**** O ******OCULAR THERAPEUTIX**

* OCULAR THERAPEUTIX INC
DEXTENZA, DEXAMETHASONE

OHM LABS

* OHM LABORATORIES INC
ACETAMINOPHEN, ACETAMINOPHEN (OTC)
IBUPROHM COLD AND SINUS, IBUPROFEN (OTC)
LOPERAMIDE HYDROCHLORIDE, LOPERAMIDE HYDROCHLORIDE (OTC)

OHM LABS INC

* OHM LABORATORIES INC
EZETIMIBE, EZETIMIBE
GUAIFENESIN, GUAIFENESIN (OTC)
VALSARTAN, VALSARTAN

OMNIVIUM PHARMS

* OMNIVIUM PHARMACEUTICALS LLC
GLYCOPYRROLATE, GLYCOPYRROLATE
ISONIAZID, ISONIAZID
MONOKET, ISOSORBIDE MONONITRATE
NUMBRINO, COCAINE HYDROCHLORIDE
SODIUM BICARBONATE, SODIUM BICARBONATE

OMSAV PHARMA

* OMSAV PHARMA RESEARCH PRIVATE LTD
NITROFURANTOIN (MONOHYDRATE/MACROCRYSTALS), NITROFURANTOIN

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** O **

ON TARGET LABS

* ON TARGET LABORATORIES INC
CYTALUX, PAFOLACIANINE SODIUM

ONYX PHARMS AMGEN

* ONYX PHARMACEUTICALS INC A WHOLLY OWNED SUB OF AMGEN INC
KYPROLIS, CARFILZOMIB

OPTINOSE US INC

* OPTINOSE US INC
XHANCE, FLUTICASONE PROPIONATE

ORAPHARMA

* ORAPHARMA INC
ARESTIN, MINOCYCLINE HYDROCHLORIDE

ORASIS PHARMS

* ORASIS PHARMACEUTICALS LTD
QLOSI, PILOCARPINE HYDROCHLORIDE

ORBICULAR

* ORBICULAR PHARMACEUTICAL TECHNOLOGIES PRIVATE LTD
ARSENIC TRIOXIDE, ARSENIC TRIOXIDE

ORBION PHARMS

* ORBION PHARMACEUTICALS PRIVATE LTD
AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
ARIPIPIRAZOLE, ARIPIPIRAZOLE
CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
CETIRIZINE HYDROCHLORIDE HIVES, CETIRIZINE HYDROCHLORIDE (OTC)
DESLORATADINE, DESLORATADINE
DIVALPROEX SODIUM, DIVALPROEX SODIUM
ESZOPICLONE, ESZOPICLONE
FELODIPINE, FELODIPINE
GRANISETRON HYDROCHLORIDE, GRANISETRON HYDROCHLORIDE
IBANDRONATE SODIUM, IBANDRONATE SODIUM
LEVETIRACETAM, LEVETIRACETAM
LEVOFLOXACIN, LEVOFLOXACIN
MODAFINIL, MODAFINIL
NARATRIPTAN, NARATRIPTAN HYDROCHLORIDE
OLANZAPINE, OLANZAPINE
PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
RASAGILINE MESYLATE, RASAGILINE MESYLATE
RISEDRONATE SODIUM, RISEDRONATE SODIUM
RIVASTIGMINE TARTRATE, RIVASTIGMINE TARTRATE
ROPINIROLE HYDROCHLORIDE, ROPINIROLE HYDROCHLORIDE
SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
TERBINAFINE HYDROCHLORIDE, TERBINAFINE HYDROCHLORIDE
VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE
ZALEPLON, ZALEPLON
ZOLMITRIPTAN, ZOLMITRIPTAN

OREXO US INC

* OREXO US INC
ZUBSOLV, BUPRENORPHINE HYDROCHLORIDE

ORGANON

* ORGANON LLC A SUB OF ORGANON AND CO
CELESTONE SOLUSPAN, BETAMETHASONE ACETATE
CLARINEX, DESLORATADINE
COZAAR, LOSARTAN POTASSIUM
DIPROLENE, BETAMETHASONE DIPROPIONATE
FOSAMAX, ALENDRONATE SODIUM
HYZAAR, HYDROCHLOROTHIAZIDE
MAXALT-MLT, RIZATRIPTAN BENZOATE
PROPECIA, FINASTERIDE
PROSCAR, FINASTERIDE
SINEMET, CARBIDOPA
SINGULAIR, MONTELUKAST SODIUM
VYTORIN, EZETIMIBE
ZETIA, EZETIMIBE
ZOCOR, SIMVASTATIN
* ORGANON USA LLC

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** O **

* ORGANON USA LLC

NEXPLANON, ETNOGESTREL
REMERON, MIRTAZAPINE

ORGANON LLC

* ORGANON LLC

ASMANEX HFA, MOMETASONE FUROATE
ASMANEX TWISTHALER, MOMETASONE FUROATE
CLARINEX-D 12 HOUR, DESLORATADINE
DULERA, FORMOTEROL FUMARATE
FOSAMAX PLUS D, ALENDRONATE SODIUM
MAXALT, RIZATRIPTAN BENZOATE
XACIATO, CLINDAMYCIN PHOSPHATE

ORGANON USA ORGANON

* ORGANON USA LLC A SUB OF ORGANON AND CO

GANIRELIX ACETATE, GANIRELIX ACETATE
NUVARING, ETHINYL ESTRADIOL
REMERON SOLTAB, MIRTAZAPINE

ORIENT PHARMA

* ORIENT PHARMA CO LTD

EZETIMIBE, EZETIMIBE

ORIENT PHARMA CO LTD

* ORIENT PHARMA CO LTD

CARISOPRODOL, CARISOPRODOL
GLYBURIDE, GLYBURIDE
PITAVASTATIN CALCIUM, PITAVASTATIN CALCIUM
VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE

ORION PHARMA

* ORION PHARMA

COMTAN, ENTACAPONE
STALEVO 100, CARBIDOPA
STALEVO 125, CARBIDOPA
STALEVO 150, CARBIDOPA
STALEVO 200, CARBIDOPA
STALEVO 50, CARBIDOPA
STALEVO 75, CARBIDOPA

ORPHALAN

* ORPHALAN SA

CUVRIOR, TRIENTINE TETRAHYDROCHLORIDE

ORYZA

* ORYZA PHARMACEUTICALS INC

DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE
OSELTAMIVIR PHOSPHATE, OSELTAMIVIR PHOSPHATE

OSMOTICA PHARM US

* OSMOTICA PHARMACEUTICAL US LLC

CARBAMAZEPINE, CARBAMAZEPINE
HYDROMORPHONE HYDROCHLORIDE, HYDROMORPHONE HYDROCHLORIDE
METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
NIFEDIPINE, NIFEDIPINE
OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE
RELEXXII, METHYLPHENIDATE HYDROCHLORIDE
VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE

OTSUKA

* OTSUKA PHARMACEUTICAL CO LTD

ABILIFY ASIMTUFII, ARIPIPRAZOLE
ABILIFY, ARIPIPRAZOLE
INQOVI, CEDAZURIDINE
JYNARQUE, TOLVAPTAN
REXULTI, BREXPIPRAZOLE
SAMSCA, TOLVAPTAN

OTSUKA PHARM

* OTSUKA PHARMACEUTICAL CO LTD

BUSULFEX, BUSULFAN

OTSUKA PHARM CO LTD

* OTSUKA PHARMACEUTICAL CO LTD

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** O ****

* OTSUKA PHARMACEUTICAL CO LTD
ABILIFY MAINTENA KIT, ARIPIPIRAZOLE

OTTER PHARMS

* OTTER PHARMACEUTICALS LLC
OTREXUP, METHOTREXATE
SYMPAZAN, CLOBAZAM

OVERSEAS

* OVERSEAS PHARMACEUTICALS LTD
LEVETIRACETAM, LEVETIRACETAM

OXFORD PHARMS

* OXFORD PHARMACEUTICALS LLC
AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
BACLOFEN, BACLOFEN
BUSPIRONE HYDROCHLORIDE, BUSPIRONE HYDROCHLORIDE
CARISOPRODOL, CARISOPRODOL
CYCLOBENZAPRINE HYDROCHLORIDE, CYCLOBENZAPRINE HYDROCHLORIDE
DIPYRIDAMOLE, DIPYRIDAMOLE
GLYCOPYRROLATE, GLYCOPYRROLATE
HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
IMIPRAMINE HYDROCHLORIDE, IMIPRAMINE HYDROCHLORIDE
LEVETIRACETAM, LEVETIRACETAM
LORAZEPAM, LORAZEPAM
METHOCARBAMOL, METHOCARBAMOL
METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
PAROXETINE HYDROCHLORIDE, PAROXETINE HYDROCHLORIDE
PRIMIDONE, PRIMIDONE
RIMACTANE, RIFAMPIN
SERTRALINE HYDROCHLORIDE, SERTRALINE HYDROCHLORIDE
SIMVASTATIN, SIMVASTATIN
SOTALOL HYDROCHLORIDE, SOTALOL HYDROCHLORIDE
SPIRONOLACTONE, SPIRONOLACTONE
TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE
TRAZODONE HYDROCHLORIDE, TRAZODONE HYDROCHLORIDE

OYSTER POINT PHARMA

* OYSTER POINT PHARMA INC
TYRVAYA, VARENICLINE TARTRATE

**** P ******P AND L**

* P AND L DEVELOPMENT LLC
ADAPALENE, ADAPALENE (OTC)
CHILDREN'S FEXOFENADINE HYDROCHLORIDE ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
CHILDREN'S FEXOFENADINE HYDROCHLORIDE HIVES, FEXOFENADINE HYDROCHLORIDE (OTC)
CLOTRIMAZOLE, CLOTRIMAZOLE (OTC)
DOCOSANOL, DOCOSANOL (OTC)
FAMOTIDINE, FAMOTIDINE (OTC)
IBUPROFEN, IBUPROFEN (OTC)
LORATADINE AND PSEUDOEPHEDRINE SULFATE, LORATADINE (OTC)
MICONAZOLE 7, MICONAZOLE NITRATE (OTC)
MINOXIDIL (FOR MEN), MINOXIDIL (OTC)
MINOXIDIL (FOR WOMEN), MINOXIDIL (OTC)
MINOXIDIL EXTRA STRENGTH (FOR MEN), MINOXIDIL (OTC)
NICOTINE POLACRILEX, NICOTINE POLACRILEX (OTC)
OMEPRAZOLE MAGNESIUM, OMEPRAZOLE MAGNESIUM (OTC)

P AND L DEV LLC

* P AND L DEVELOPMENT LLC DBA PLD DEVELOPMENTS LLC
IBUPROFEN, IBUPROFEN (OTC)

P AND L DEVELOPMENT

* P AND L DEVELOPMENT LLC
DOCOSANOL, DOCOSANOL (OTC)

PACIFIC PHARMA

* PACIFIC PHARMA
TIMOLOL MALEATE, TIMOLOL MALEATE
* PACIFIC PHARMA INC
TIMOLOL MALEATE, TIMOLOL MALEATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** P **

PACIRA PHARMS INC

* PACIRA PHARMACEUTICALS INC
 EXPAREL, BUPIVACAINE
 ZILRETTA, TRIAMCINOLONE ACETONIDE

PADAGIS ISRAEL

* PADAGIS ISRAEL PHARMACEUTICALS LTD
 ACYCLOVIR, ACYCLOVIR
 ADAPALENE AND BENZOYL PEROXIDE, ADAPALENE
 AMMONIUM LACTATE, AMMONIUM LACTATE
 AZELASTINE HYDROCHLORIDE AND FLUTICASONE PROPIONATE, AZELASTINE HYDROCHLORIDE
 BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
 BETAMETHASONE VALERATE, BETAMETHASONE VALERATE
 BRIMONIDINE TARTRATE, BRIMONIDINE TARTRATE
 BUDESONIDE, BUDESONIDE
 CALCIPOTRIENE AND BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
 CALCIPOTRIENE AND BETHAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
 CICLOPIROX, CICLOPIROX
 CLINDAMYCIN PHOSPHATE AND BENZOYL PEROXIDE, BENZOYL PEROXIDE
 CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
 CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
 CYANOCOBALAMIN, CYANOCOBALAMIN
 DESOXIMETASONE, DESOXIMETASONE
 DICLOFENAC SODIUM, DICLOFENAC SODIUM
 ECONAZOLE NITRATE, ECONAZOLE NITRATE
 ESTRADIOL, ESTRADIOL
 FLUOCINOLONE ACETONIDE, FLUOCINOLONE ACETONIDE
 FLURANDRENOLIDE, FLURANDRENOLIDE
 FLUTICASONE PROPIONATE, FLUTICASONE PROPIONATE
 GYNAZOLE-1, BUTOCONAZOLE NITRATE
 HALOBETASOL PROPIONATE, HALOBETASOL PROPIONATE
 HYDROCORTISONE VALERATE, HYDROCORTISONE VALERATE
 IMIQUIMOD, IMIQUIMOD
 IVERMECTIN, IVERMECTIN
 KETOCONAZOLE, KETOCONAZOLE
 MESALAMINE, MESALAMINE
 METRONIDAZOLE, METRONIDAZOLE
 MOMETASONE FUROATE, MOMETASONE FUROATE
 MUPIROCIN, MUPIROCIN
 NALOXONE HYDROCHLORIDE, NALOXONE HYDROCHLORIDE (OTC)
 NITROGLYCERIN, NITROGLYCERIN
 NYSTATIN AND TRIAMCINOLONE ACETONIDE, NYSTATIN
 OLOPATADINE HYDROCHLORIDE, OLOPATADINE HYDROCHLORIDE
 PENCICLOVIR, PENCICLOVIR
 PERMETHRIN, PERMETHRIN
 PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
 SUMATRIPTAN, SUMATRIPTAN
 TAZAROTENE, TAZAROTENE
 TERCONAZOLE, TERCONAZOLE
 TESTOSTERONE, TESTOSTERONE
 TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE
 ZOLMITRIPTAN, ZOLMITRIPTAN

PADAGIS US

* PADAGIS US LLC
 BACITRACIN ZINC AND POLYMYXIN B SULFATE, BACITRACIN ZINC
 BACITRACIN, BACITRACIN
 BACITRACIN-NEOMYCIN-POLYMYXIN W/ HYDROCORTISONE ACETATE, BACITRACIN ZINC
 BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
 BRINZOLAMIDE, BRINZOLAMIDE
 BROMOCRIPTINE MESYLATE, BROMOCRIPTINE MESYLATE
 BROMPHENIRAMINE MALEATE, PSEUDOEPHEDRINE HYDROCHLORIDE AND DEXTROMETHORPHAN
 CALCIUM ACETATE, CALCIUM ACETATE
 CENTANY, MUPIROCIN
 CETIRIZINE HYDROCHLORIDE, CETIRIZINE HYDROCHLORIDE
 CICLOPIROX, CICLOPIROX
 CLINDA-DERM, CLINDAMYCIN PHOSPHATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** P ******* PADAGIS US LLC**

CLINDAMYCIN PALMITATE HYDROCHLORIDE, CLINDAMYCIN PALMITATE HYDROCHLORIDE
 CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
 CLINDESSE, CLINDAMYCIN PHOSPHATE
 CLINDETS, CLINDAMYCIN PHOSPHATE
 CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
 CLOTRIMAZOLE, CLOTRIMAZOLE
 COMPRO, PROCHLORPERAZINE
 CYCLOSPORINE, CYCLOSPORINE
 DESONIDE, DESONIDE
 DESOXIMETASONE, DESOXIMETASONE
 DIHYDROERGOTAMINE MESYLATE, DIHYDROERGOTAMINE MESYLATE
 ENTOCORT EC, BUDESONIDE
 ERYTHROMYCIN, ERYTHROMYCIN
 EVAMIST, ESTRADIOL
 FLAVOXATE HYDROCHLORIDE, FLAVOXATE HYDROCHLORIDE
 GENTAMICIN SULFATE, GENTAMICIN SULFATE
 HYDROCODONE BITARTRATE AND HOMATROPINE METHYLBROMIDE, HOMATROPINE METHYLBROMIDE
 HYDROCODONE BITARTRATE, CHLORPHENIRAMINE MALEATE AND PSEUDOEPHEDRINE HYDROCHLORIDE,
 HYDROCORTISONE, HYDROCORTISONE
 HYDROMORPHONE HYDROCHLORIDE, HYDROMORPHONE HYDROCHLORIDE
 IBUPROFEN, IBUPROFEN
 LEVOCETIRIZINE DIHYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE
 LIDOCAINE AND PRILOCAINE, LIDOCAINE
 LOTE Prednol Etabonate, LOTE Prednol Etabonate
 MIDAMOR, AMILORIDE HYDROCHLORIDE
 MIDAZOLAM HYDROCHLORIDE, MIDAZOLAM HYDROCHLORIDE
 MOMETASONE FUROATE, MOMETASONE FUROATE
 MORPHINE SULFATE, MORPHINE SULFATE
 NARATRIPTAN, NARATRIPTAN HYDROCHLORIDE
 NEOMYCIN AND POLYMYXIN B SULFATES AND BACITRACIN ZINC, BACITRACIN ZINC
 NEOMYCIN AND POLYMYXIN B SULFATES AND DEXAMETHASONE, DEXAMETHASONE
 NYSTATIN, NYSTATIN
 NYSTOP, NYSTATIN
 PILOCARPINE HYDROCHLORIDE, PILOCARPINE HYDROCHLORIDE
 PODOFILOX, PODOFILOX
 POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
 REPAGLINIDE, REPAGLINIDE
 SCOPOLAMINE, SCOPOLAMINE
 SELENIUM SULFIDE, SELENIUM SULFIDE
 STIE-CORT, HYDROCORTISONE
 SULFACETAMIDE SODIUM, SULFACETAMIDE SODIUM
 TAVABOROLE, TAVABOROLE
 TESTOSTERONE CYPIONATE, TESTOSTERONE CYPIONATE
 TOBRAMYCIN AND DEXAMETHASONE, DEXAMETHASONE
 TRETINOIN, TRETINOIN
 TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE
 TROSPIMUM CHLORIDE, TROSPIMUM CHLORIDE

PAI HOLDINGS

* PAI HOLDINGS LLC DBA PHARMACEUTICAL ASSOCIATES INC AND DBA PAI PHARMA
 SODIUM SULFATE, POTASSIUM SULFATE AND MAGNESIUM SULFATE, MAGNESIUM SULFATE

PAI HOLDINGS PHARM

* PAI HOLDINGS LLC DBA PHARMACEUTICAL ASSOCIATES INC
 CEFOTAN, CEFOTETAN DISODIUM
 CIMETIDINE HYDROCHLORIDE, CIMETIDINE HYDROCHLORIDE
 HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE
 IBUPROFEN, IBUPROFEN (OTC)
 LIDOCAINE AND PRILOCAINE, LIDOCAINE
 LIDOCAINE HYDROCHLORIDE VISCOUS, LIDOCAINE HYDROCHLORIDE
 LIDOCAINE HYDROCHLORIDE, LIDOCAINE HYDROCHLORIDE

PANACEA

* PANACEA BIOTEC PHARMA LTD
 PRASUGREL, PRASUGREL HYDROCHLORIDE
 RIZATRIPTAN BENZOATE, RIZATRIPTAN BENZOATE
 TACROLIMUS, TACROLIMUS

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** P **

PANGEA

- * PANGEA PHARMACEUTICALS LLC
ERGOMAR, ERGOTAMINE TARTRATE
ETODOLAC, ETODOLAC
OXAPROZIN, OXAPROZIN

PARAGON BIOTECK

- * PARAGON BIOTECK INC
PHENYLEPHRINE HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE

PARATEK PHARMS INC

- * PARATEK PHARMACEUTICALS INC
NUZYRA, OMADACYCLINE TOSYLATE

PARKE DAVIS

- * PARKE DAVIS DIV WARNER LAMBERT CO
CELONTIN, METHSUXIMIDE
CEREBYX, FOSPHENYTOIN SODIUM
NARDIL, PHENELZINE SULFATE
ZARONTIN, ETHOSUXIMIDE

PARKE-DAVIS

- * PARKE-DAVIS DIVISION OF PFIZER INC
ZARONTIN, ETHOSUXIMIDE

PATHEON SOFTGELS

- * PATHEON SOFTGELS BV
IBUPROFEN, IBUPROFEN (OTC)
NAPROXEN SODIUM, NAPROXEN SODIUM (OTC)

PD PARTNERS

- * PD PARTNERS III LLC
SUCRALFATE, SUCRALFATE

PENN LIFE

- * PENN LIFE SCIENCES LLC
ARSENIC TRIOXIDE, ARSENIC TRIOXIDE
CARMUSTINE, CARMUSTINE
ISOPROTERENOL HYDROCHLORIDE, ISOPROTERENOL HYDROCHLORIDE
THIOTEPA, THIOTEPA

PERRIGO NEW YORK

- * PERRIGO NEW YORK INC
ACETAMINOPHEN, ACETAMINOPHEN (OTC)
PERMETHRIN, PERMETHRIN (OTC)

PERRIGO PHARMA INTL

- * PERRIGO PHARMA INTERNATIONAL DAC
DICLOFENAC SODIUM, DICLOFENAC SODIUM (OTC)
LORATADINE AND PSEUDOEPHEDRINE SULFATE, LORATADINE (OTC)
LORATADINE, LORATADINE (OTC)
MINOXIDIL, MINOXIDIL (OTC)
NASONEX 24HR ALLERGY, MOMETASONE FUROATE (OTC)
TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE (OTC)
- * PERRIGO PHARMA INTERNATIONAL DESIGNATED ACTIVITY CO
LORATADINE, LORATADINE (OTC)
PREVACID 24 HR, LANSOPRAZOLE (OTC)

PERRIGO R AND D

- * PERRIGO R AND D CO
CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
CETIRIZINE HYDROCHLORIDE HIVES, CETIRIZINE HYDROCHLORIDE (OTC)
CHILDREN'S CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
CHILDREN'S CETIRIZINE HYDROCHLORIDE HIVES RELIEF, CETIRIZINE HYDROCHLORIDE (OTC)
ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM (OTC)
FAMOTIDINE, CALCIUM CARBONATE, AND MAGNESIUM HYDROXIDE, CALCIUM CARBONATE (OTC)
FAMOTIDINE, FAMOTIDINE (OTC)
GUAIFENESIN AND DEXTROMETHORPHAN HYDROBROMIDE, DEXTROMETHORPHAN HYDROBROMIDE (OTC)
GUAIFENESIN, GUAIFENESIN (OTC)
IBUPROFEN AND DIPHENHYDRAMINE CITRATE, DIPHENHYDRAMINE CITRATE (OTC)
IBUPROFEN AND PHENYLEPHRINE HYDROCHLORIDE, IBUPROFEN (OTC)
IBUPROFEN, IBUPROFEN (OTC)
LANSOPRAZOLE, LANSOPRAZOLE (OTC)
LEVOCETIRIZINE DIHYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE (OTC)

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** P ****

- * PERRIGO R AND D CO
 - LEVONORGESTREL, LEVONORGESTREL (OTC)
 - LOPERAMIDE HYDROCHLORIDE AND SIMETHICONE, LOPERAMIDE HYDROCHLORIDE (OTC)
 - LOPERAMIDE HYDROCHLORIDE, LOPERAMIDE HYDROCHLORIDE (OTC)
 - MICONAZOLE NITRATE, MICONAZOLE NITRATE (OTC)
 - NAPROXEN SODIUM AND DIPHENHYDRAMINE HYDROCHLORIDE, DIPHENHYDRAMINE HYDROCHLORIDE
 - NICOTINE POLACRILEX, NICOTINE POLACRILEX (OTC)
 - OMEPRAZOLE AND SODIUM BICARBONATE, OMEPRAZOLE (OTC)

PETNET

- * PETNET SOLUTIONS INC
 - AMMONIA N 13, AMMONIA N-13
 - FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18
 - SODIUM FLUORIDE F-18, SODIUM FLUORIDE F-18

PF PRISM CV

- * PF PRISM CV
 - BOSULIF, BOSUTINIB MONOHYDRATE
 - INLYTA, AXITINIB
 - PRISTIQ, DESVENLAFAXINE SUCCINATE
 - RAPAMUNE, SIROLIMUS
 - TORISEL, TEMSIROLIMUS
 - TYGACIL, TIGECYCLINE
 - VFEND, VORICONAZOLE
 - XALKORI, CRIZOTINIB
 - XELJANZ, TOFACITINIB CITRATE

PFIZER

- * PFIZER CENTRAL RESEARCH
 - DIFLUCAN, FLUCONAZOLE
 - ZITHROMAX, AZITHROMYCIN
- * PFIZER CHEMICALS DIV PFIZER INC
 - DIFLUCAN, FLUCONAZOLE
 - ZITHROMAX, AZITHROMYCIN
- * PFIZER INC
 - ALDACTAZIDE, HYDROCHLOROTHIAZIDE
 - ALDACTONE, SPIRONOLACTONE
 - AROMASIN, EXEMESTANE
 - ARTHROTEC, DICLOFENAC SODIUM
 - AZULFIDINE EN-TABS, SULFASALAZINE
 - AZULFIDINE, SULFASALAZINE
 - CAVERJECT IMPULSE, ALPROSTADIL
 - CAVERJECT, ALPROSTADIL
 - CIBINQO, ABROCITINIB
 - CLEOCIN HYDROCHLORIDE, CLINDAMYCIN HYDROCHLORIDE
 - CLEOCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
 - CLEOCIN T, CLINDAMYCIN PHOSPHATE
 - CLEOCIN, CLINDAMYCIN PALMITATE HYDROCHLORIDE
 - CLEOCIN, CLINDAMYCIN PHOSPHATE
 - COLESTID, COLESTIPOL HYDROCHLORIDE
 - CORVERT, IBUTILIDE FUMARATE
 - CYKLOKAPRON, TRANEXAMIC ACID
 - CYTOTEC, MISOPROSTOL
 - DAURISMO, GLASDEGIB MALEATE
 - DAYPRO, OXAPROZIN
 - DEPO-ESTRADIOL, ESTRADIOL CYPIONATE
 - DEPO-MEDROL, METHYLPREDNISOLONE ACETATE
 - DEPO-PROVERA, MEDROXYPROGESTERONE ACETATE
 - DEPO-SUBQ PROVERA 104, MEDROXYPROGESTERONE ACETATE
 - DEPO-TESTOSTERONE, TESTOSTERONE CYPIONATE
 - DOXORUBICIN HYDROCHLORIDE, DOXORUBICIN HYDROCHLORIDE
 - ESTRING, ESTRADIOL
 - FLAGYL, METRONIDAZOLE
 - FRAGMIN, DALTEPARIN SODIUM
 - GLUCOTROL XL, GLIPIZIDE
 - GLYNASE, GLYBURIDE
 - GLYSET, MIGLITOL
 - HALCION, TRIAZOLAM

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** P ****

- * PFIZER INC
 HEMABATE, CARBOPROST TROMETHAMINE
 HEPARIN SODIUM PRESERVATIVE FREE, HEPARIN SODIUM
 HEPARIN SODIUM, HEPARIN SODIUM
 IBRANCE, PALBOCICLIB
 IDAMYCIN PFS, IDARUBICIN HYDROCHLORIDE
 LINCOCIN, LINCOMYCIN HYDROCHLORIDE
 LITFULO, RITLECITINIB TOSYLATE
 LOMOTIL, ATROPINE SULFATE
 LORBRENA, LORLATINIB
 MEDROL, METHYLPREDNISOLONE
 MERREM IV, MEROPENEM
 MYCOBUTIN, RIFABUTIN
 NORPACE CR, DISOPYRAMIDE PHOSPHATE
 NORPACE, DISOPYRAMIDE PHOSPHATE
 NURTEC ODT, RIMEGEPANT SULFATE
 OGEN 5, ESTROPIPATE
 PAXLOVID (COPACKAGED), NIRMATRELVIR
 PREPIDIL, DINOPROSTONE
 PROCARDIA, NIFEDIPINE
 PROSTIN VR PEDIATRIC, ALPROSTADIL
 PROVERA, MEDROXYPROGESTERONE ACETATE
 SONATA, ZALEPLON
 SYNAREL, NAFARELIN ACETATE
 TALZENNA, TALAZOPARIB TOSYLATE
 TESSALON, BENZONATATE
 TOVIAZ, FESOTERODINE FUMARATE
 UNASYN, AMPICILLIN SODIUM
 VELSIPITY, ETRASIMOD ARGININE
 VIZIMPRO, DACOMITINIB
 XELJANZ XR, TOFACITINIB CITRATE
 XELJANZ, TOFACITINIB CITRATE
 ZAVZPRET, ZAVEGEPANT HYDROCHLORIDE
 ZITHROMAX, AZITHROMYCIN
 ZYVOX, LINEZOLID
- * PFIZER LABORATORIES DIV PFIZER INC
 FELDENE, PIROXICAM
 MINIPRESS, PRAZOSIN HYDROCHLORIDE
 PFIZERPEN, PENICILLIN G POTASSIUM
 PROCARDIA XL, NIFEDIPINE
 VIBRAMYCIN, DOXYCYCLINE HYCLATE
 VISTARIL, HYDROXYZINE PAMOATE
- * PFIZER PHARMACEUTICALS PRODUCTION CORP LTD
 TIKOSYN, DOFETILIDE
- PFIZER INC**
- * PFIZER INC
 CAMPTOSAR, IRINOTECAN HYDROCHLORIDE
 ELLENCE, EPIRUBICIN HYDROCHLORIDE
 NICOTROL, NICOTINE
- PFIZER PHARMS**
- * PFIZER PHARMACEUTICALS LTD
 LOPID, GEMFIBROZIL
- PHARM ASSOC**
- * PHARMACEUTICAL ASSOC INC
 HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
 LORAZEPAM, LORAZEPAM
 PREDNISOLONE SODIUM PHOSPHATE, PREDNISOLONE SODIUM PHOSPHATE
- * PHARMACEUTICAL ASSOCIATES INC
 AMANTADINE HYDROCHLORIDE, AMANTADINE HYDROCHLORIDE
 BROMPHENIRAMINE MALEATE, PSEUDOEPHEDRINE HYDROCHLORIDE AND DEXTROMETHORPHAN
 CETIRIZINE HYDROCHLORIDE, CETIRIZINE HYDROCHLORIDE
 CHLORHEXIDINE GLUCONATE, CHLORHEXIDINE GLUCONATE
 DIPHENHYDRAMINE HYDROCHLORIDE, DIPHENHYDRAMINE HYDROCHLORIDE
 ETHOSUXIMIDE, ETHOSUXIMIDE
 FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** P ****

* PHARMACEUTICAL ASSOCIATES INC
 FLUPHENAZINE HYDROCHLORIDE, FLUPHENAZINE HYDROCHLORIDE
 HALOPERIDOL, HALOPERIDOL LACTATE
 HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
 LACTULOSE, LACTULOSE
 LEVETIRACETAM, LEVETIRACETAM
 NORTRIPTYLINE HYDROCHLORIDE, NORTRIPTYLINE HYDROCHLORIDE
 NYSTATIN, NYSTATIN
 OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
 POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
 PREDNISOLONE, PREDNISOLONE
 PROMETHAZINE WITH CODEINE, CODEINE PHOSPHATE
 SULFATRIM PEDIATRIC, SULFAMETHOXAZOLE
 THEOPHYLLINE, THEOPHYLLINE
 TRIHEXYPHENIDYL HYDROCHLORIDE, TRIHEXYPHENIDYL HYDROCHLORIDE
 VALPROIC ACID, VALPROIC ACID

PHARMAAND

* PHARMAAND GMBH
 RUBRACA, RUCAPARIB CAMSYLATE

PHARMACHEMIE BV

* PHARMACHEMIE BV
 CARBOPLATIN, CARBOPLATIN
 CISPLATIN, CISPLATIN
 METHOTREXATE SODIUM PRESERVATIVE FREE, METHOTREXATE SODIUM

PHARMACIA

* PHARMACIA AND UPJOHN CO LLC
 CADUET, AMLODIPINE BESYLATE
 DILANTIN, PHENYTOIN

PHARMACIA AND UPJOHN

* PHARMACIA AND UPJOHN CO
 CORTEF, HYDROCORTISONE
 R-GENE 10, ARGININE HYDROCHLORIDE
 SOLU-CORTEF, HYDROCORTISONE SODIUM SUCCINATE
 SOLU-MEDROL, METHYLPREDNISOLONE SODIUM SUCCINATE

PHARMACOSMOS

* PHARMACOSMOS AS
 COSELA, TRILACICLIB DIHYDROCHLORIDE
 MONOFERRIC, FERRIC DERISOMALTOSE

PHARMACYCLICS LLC

* PHARMACYCLICS LLC
 IMBRUVICA, IBRUTINIB

PHARMADAX INC

* PHARMADAX INC
 METOPROLOL SUCCINATE, METOPROLOL SUCCINATE

PHARMALOGIC

* PHARMALOGIC SOUTH CAROLINA LLC
 FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18

PHARMALOGIC HLDGS

* PHARMALOGIC HOLDINGS CORP
 FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18

PHARMASCIENCE INC

* PHARMASCIENCE INC
 BORTEZOMIB, BORTEZOMIB
 BUSULFAN, BUSULFAN
 DECITABINE, DECITABINE
 GANCICLOVIR SODIUM, GANCICLOVIR SODIUM

PHARMAXIS EUROPE

* PHARMAXIS EUROPE LTD
 ARIDOL KIT, MANNITOL
 BRONCHITOL, MANNITOL

PHARMING

* PHARMING TECHNOLOGIES BV
 JOENJA, LENIOLISIB PHOSPHATE

PHARMOBEDIENT

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** P ****

* PHARMOBEDIENT CONSULTING LLC
CHLORTHALIDONE, CHLORTHALIDONE

PHATHOM

* PHATHOM PHARMACEUTICALS INC
VOQUEZNA DUAL PAK, AMOXICILLIN
VOQUEZNA TRIPLE PAK, AMOXICILLIN
VOQUEZNA, VONOPRAZAN FUMARATE

PHOTOCURE ASA

* PHOTOCURE ASA
CYSVIEW KIT, HEXAMINOLEVULINATE HYDROCHLORIDE

PIERRE FABRE

* PIERRE FABRE MEDICAMENT
HEMANGEOL, PROPRANOLOL HYDROCHLORIDE

PIERREL

* PIERREL S.P.A.
ORABLOC, ARTICAINA HYDROCHLORIDE

PINNACLE BIOLGS

* PINNACLE BIOLOGICS INC
PHOTOFRIN, PORFIMER SODIUM

PIRAMAL

* PIRAMAL HEALTHCARE UK LTD
CLOBAZAM, CLOBAZAM
DEFERASIROX, DEFERASIROX
VARENICLINE TARTRATE, VARENICLINE TARTRATE

PIRAMAL CRITICAL

* PIRAMAL CRITICAL CARE INC
CISATRACURIUM BESYLATE, CISATRACURIUM BESYLATE
FOSAPREPITANT DIMEGLUMINE, FOSAPREPITANT DIMEGLUMINE
GABLOFEN, BACLOFEN
GLYCOPYRROLATE, GLYCOPYRROLATE
ISOFLURANE, ISOFLURANE
MITIGO, MORPHINE SULFATE
ROCURONIUM BROMIDE, ROCURONIUM BROMIDE
SOJOURN, SEVOFLURANE
* PIRAMAL CRITICAL CARE LTD
LEVOTHYROXINE SODIUM, LEVOTHYROXINE SODIUM

PIRAMAL PHARMA

* PIRAMAL PHARMA LTD
ISOFLURANE, ISOFLURANE

PLD ACQUISITIONS

* PLD ACQUISITIONS LLC DBA AVEMA PHARMA SOLUTIONS
CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
CETIRIZINE HYDROCHLORIDE AND PSEUDOEPHEDRINE HYDROCHLORIDE, CETIRIZINE HYDROCHLORIDE
NICOTINE POLACRILEX, NICOTINE POLACRILEX (OTC)

PLD ACQUISITIONS LLC

* PLD ACQUISITIONS LLC
IBUPROFEN AND DIPHENHYDRAMINE CITRATE, DIPHENHYDRAMINE CITRATE (OTC)
LORATADINE, LORATADINE (OTC)
ZOLMITRIPTAN, ZOLMITRIPTAN

PLIVA

* PLIVA INC
AZITHROMYCIN, AZITHROMYCIN
BENZTROPINE MESYLATE, BENZTROPINE MESYLATE
HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE
METRONIDAZOLE, METRONIDAZOLE
TRAZODONE HYDROCHLORIDE, TRAZODONE HYDROCHLORIDE
WARFARIN SODIUM, WARFARIN SODIUM

PLIVA PHARM IND

* PLIVA PHARMACEUTICAL INDUSTRY INC
TORSEMIDE, TORSEMIDE

PLX PHARMA

* PLX PHARMA INC
VAZALORE, ASPIRIN (OTC)

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** P ******POHL BOSKAMP**

* POHL BOSKAMP
NITROLINGUAL PUMPSPRAY, NITROGLYCERIN

POLAREAN

* POLAREAN INC
XENOVIEW, XENON XE-129 HYPERPOLARIZED

POLYGEN PHARMS

* POLYGEN PHARMACEUTICALS INC
AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
CLOPIDOGREL BISULFATE, CLOPIDOGREL BISULFATE
DARIFENACIN HYDROBROMIDE, DARIFENACIN HYDROBROMIDE
MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE

POWDER PHARMS

* POWDER PHARMACEUTICALS INC
ZINGO, LIDOCAINE HYDROCHLORIDE

PPI-DAC

* PERRIGO PHARMA INTERNATIONAL DAC
CETIRIZINE HYDROCHLORIDE AND PSEUDOEPHEDRINE HYDROCHLORIDE, CETIRIZINE HYDROCHLORIDE
NICOTINE POLACRILEX, NICOTINE POLACRILEX (OTC)
POLYETHYLENE GLYCOL 3350, POLYETHYLENE GLYCOL 3350 (OTC)

PRASCO

* PRASCO LLC DBA PRASCO LABORATORIES
DEXAMETHASONE, DEXAMETHASONE
DEXTROAMPHETAMINE SULFATE, DEXTROAMPHETAMINE SULFATE
ESTRADIOL, ESTRADIOL
MIRTAZAPINE, MIRTAZAPINE
SULFAMETHOXAZOLE AND TRIMETHOPRIM, SULFAMETHOXAZOLE

PRAXGEN

* PRAXGEN PHARMACEUTICALS LLC
RANOLAZINE, RANOLAZINE

PRECISION DERMAT

* PRECISION DERMATOLOGY INC
LOCOID LIPOCREAM, HYDROCORTISONE BUTYRATE
LOCOID, HYDROCORTISONE BUTYRATE

PRECISION DOSE INC

* PRECISION DOSE INC
PHENTOLAMINE MESYLATE, PHENTOLAMINE MESYLATE

PRECISION NUCLEAR

* PRECISION NUCLEAR LLC
AMMONIA N 13, AMMONIA N-13
FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18
SODIUM FLUORIDE F-18, SODIUM FLUORIDE F-18

PRIMUS PHARMS

* PRIMUS PHARMACEUTICALS INC
IMPOYZ, CLOBETASOL PROPIONATE
SERNIVO, BETAMETHASONE DIPROPIONATE

PRINSTON INC

* PRINSTON PHARMACEUTICAL INC
AMPHETAMINE SULFATE, AMPHETAMINE SULFATE
ARIPIRAZOLE, ARIPIRAZOLE
BENAZEPRIL HYDROCHLORIDE, BENAZEPRIL HYDROCHLORIDE
BISOPROLOL FUMARATE, BISOPROLOL FUMARATE
BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
BUSULFAN, BUSULFAN
CANDESARTAN CILEXETIL AND HYDROCHLOROTHIAZIDE, CANDESARTAN CILEXETIL
CANDESARTAN CILEXETIL, CANDESARTAN CILEXETIL
CAPTOPRIL, CAPTOPRIL
CLONAZEPAM, CLONAZEPAM
CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
CLOPIDOGREL BISULFATE, CLOPIDOGREL BISULFATE
CYCLOBENZAPRINE HYDROCHLORIDE, CYCLOBENZAPRINE HYDROCHLORIDE
DICYCLOMINE HYDROCHLORIDE, DICYCLOMINE HYDROCHLORIDE
DIMETHYL FUMARATE, DIMETHYL FUMARATE
DIVALPROEX SODIUM, DIVALPROEX SODIUM

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** P ****

* PRINSTON PHARMACEUTICAL INC
 DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
 DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
 DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE
 ENALAPRIL MALEATE, ENALAPRIL MALEATE
 ESCITALOPRAM OXALATE, ESCITALOPRAM OXALATE
 ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM
 FEBUXOSTAT, FEBUXOSTAT
 FENOFIBRATE, FENOFIBRATE
 FINGOLIMOD HYDROCHLORIDE, FINGOLIMOD HYDROCHLORIDE
 FOSINOPRIL SODIUM, FOSINOPRIL SODIUM
 FUROSEMIDE, FUROSEMIDE
 GLIMEPIRIDE, GLIMEPIRIDE
 GLYCOPYRROLATE, GLYCOPYRROLATE
 HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE
 IRBESARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 IRBESARTAN, IRBESARTAN
 LEVETIRACETAM, LEVETIRACETAM
 LEVOMILNACIPRAN HYDROCHLORIDE, LEVOMILNACIPRAN HYDROCHLORIDE
 LISDEXAMFETAMINE DIMESYLATE, LISDEXAMFETAMINE DIMESYLATE
 LISINOPRIL AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 LISINOPRIL, LISINOPRIL
 LOSARTAN POTASSIUM AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 LOSARTAN POTASSIUM, LOSARTAN POTASSIUM
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 METHOCARBAMOL, METHOCARBAMOL
 METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
 NEBIVOLOL HYDROCHLORIDE, NEBIVOLOL HYDROCHLORIDE
 OLMESARTAN MEDOXOMIL AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 OLMESARTAN MEDOXOMIL, OLMESARTAN MEDOXOMIL
 PAROXETINE HYDROCHLORIDE, PAROXETINE HYDROCHLORIDE
 PAROXETINE MESYLATE, PAROXETINE MESYLATE
 PAROXETINE, PAROXETINE HYDROCHLORIDE
 PEMETREXED DISODIUM, PEMETREXED DISODIUM
 PHENTERMINE HYDROCHLORIDE, PHENTERMINE HYDROCHLORIDE
 PIOGLITAZONE HYDROCHLORIDE, PIOGLITAZONE HYDROCHLORIDE
 POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
 PREGABALIN, PREGABALIN
 PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
 QUETIAPINE FUMARATE, QUETIAPINE FUMARATE
 QUINAPRIL HYDROCHLORIDE, QUINAPRIL HYDROCHLORIDE
 RISPERIDONE, RISPERIDONE
 ROCURONIUM BROMIDE, ROCURONIUM BROMIDE
 ROFLUMILAST, ROFLUMILAST
 ROPINIROLE HYDROCHLORIDE, ROPINIROLE HYDROCHLORIDE
 SAFINAMIDE MESYLATE, SAFINAMIDE MESYLATE
 SILODOSIN, SILODOSIN
 TADALAFIL, TADALAFIL
 TELMISARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 TELMISARTAN, TELMISARTAN
 TEMAZEPAM, TEMAZEPAM
 TICAGRELOR, TICAGRELOR
 VALSARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 VALSARTAN, VALSARTAN
 VORICONAZOLE, VORICONAZOLE
 VYDUO, NEBIVOLOL HYDROCHLORIDE

PROF DSPLS

* PROFESSIONAL DISPOSABLES INC
 PREVANTICS MAXI SWABSTICK, CHLORHEXIDINE GLUCONATE (OTC)
 PREVANTICS SWAB, CHLORHEXIDINE GLUCONATE (OTC)
 PREVANTICS SWABSTICK, CHLORHEXIDINE GLUCONATE (OTC)

PROGENICS PHARMS INC

* PROGENICS PHARMACEUTICALS INC
 PYLARIFY, PIFLUFOLASTAT F-18

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** P ******PROTEGA PHARMS**

* PROTEGA PHARMACEUTICALS INC
ROXYBOND, OXYCODONE HYDROCHLORIDE

PROVELL

* PROVELL PHARMACEUTICALS LLC
EUTHYROX, LEVOTHYROXINE SODIUM **

PROVENSIS

* PROVENSIS LTD
VARITHENA, POLIDOCANOL

PROVEPHARM SAS

* PROVEPHARM SAS
BAL, DIMERCAPROL
BLUDIGO, INDIGOTINDISULFONATE SODIUM
DIHYDROERGOTAMINE MESYLATE, DIHYDROERGOTAMINE MESYLATE
PHENYLEPHRINE HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE
PIPERACILLIN AND TAZOBACTAM, PIPERACILLIN SODIUM
PROVAYBLUE, METHYLENE BLUE
TRANEXAMIC ACID, TRANEXAMIC ACID

PTC THERAP

* PTC THERAPEUTICS INC
EMFLAZA, DEFLAZACORT

PULMOFLOW INC

* PULMOFLOW INC
KITABIS PAK, TOBRAMYCIN

PUMA BIOTECH

* PUMA BIOTECHNOLOGY INC
NERLYNX, NERATINIB MALEATE

PURACAP PHARM

* PURACAP PHARMACEUTICAL LLC
MELOXICAM, MELOXICAM

PURACAP PHARM LLC

* PURACAP PHARMACEUTICAL LLC
BENZONATATE, BENZONATATE
ERGOCALCIFEROL, ERGOCALCIFEROL
ETHOSUXIMIDE, ETHOSUXIMIDE
MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
NAPROXEN SODIUM, NAPROXEN SODIUM (OTC)
OMEGA-3-ACID ETHYL ESTERS, OMEGA-3-ACID ETHYL ESTERS
PIOGLITAZONE HYDROCHLORIDE, PIOGLITAZONE HYDROCHLORIDE

PURDUE PHARMA LP

* PURDUE PHARMA LP
BUTRANS, BUPRENORPHINE
HYSINGLA ER, HYDROCODONE BITARTRATE
MS CONTIN, MORPHINE SULFATE
NALMEFENE HYDROCHLORIDE, NALMEFENE HYDROCHLORIDE
OXYCONTIN, OXYCODONE HYDROCHLORIDE
ZURNAI (AUTOINJECTOR), NALMEFENE HYDROCHLORIDE

PURE SOURCE

* PURE SOURCE LLC
THEROXIDIL, MINOXIDIL (OTC)

PYROS PHARMS

* PYROS PHARMACEUTICALS INC
VIGAFYDE, VIGABATRIN
VIGPODER, VIGABATRIN

**** Q ******Q BIOMED**

* Q BIOMED INC
METASTRON, STRONTIUM CHLORIDE SR-89
STRONTIUM CHLORIDE SR-89, STRONTIUM CHLORIDE SR-89

QILU

* QILU PHARMACEUTICAL CO LTD
ABIRATERONE ACETATE, ABIRATERONE ACETATE
AMIKACIN SULFATE, AMIKACIN SULFATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** Q ****

* QILU PHARMACEUTICAL CO LTD
 CEFAZOLIN SODIUM, CEFAZOLIN SODIUM
 CEFEPIME HYDROCHLORIDE, CEFEPIME HYDROCHLORIDE
 CEFTRIAZONE, CEFTRIAZONE SODIUM
 CETRORELIX ACETATE, CETRORELIX ACETATE
 CISPLATIN, CISPLATIN
 EXEMESTANE, EXEMESTANE
 HYDROXYUREA, HYDROXYUREA
 ICOSAPENT ETHYL, ICOSAPENT ETHYL
 MEROPENEM, MEROPENEM
 OLANZAPINE, OLANZAPINE
 OLMESARTAN MEDOXOMIL, OLMESARTAN MEDOXOMIL
 SOLIFENACIN SUCCINATE, SOLIFENACIN SUCCINATE
 TENOFOVIR DISOPROXIL FUMARATE, TENOFOVIR DISOPROXIL FUMARATE

QILU ANTIBIOTICS

* QILU ANTIBIOTICS PHARMACEUTICAL CO LTD
 ERTAPENEM SODIUM, ERTAPENEM SODIUM

QILU PHARM HAINAN

* QILU PHARMACEUTICAL HAINAN CO LTD
 BORTEZOMIB, BORTEZOMIB
 BUMETANIDE, BUMETANIDE
 DAPTOMYCIN, DAPTOMYCIN
 DECITABINE, DECITABINE
 FOSAPREPITANT DIMEGLUMINE, FOSAPREPITANT DIMEGLUMINE
 GEFITINIB, GEFITINIB
 IMATINIB MESYLATE, IMATINIB MESYLATE
 IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE
 ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
 OXALIPLATIN, OXALIPLATIN
 PALONOSETRON HYDROCHLORIDE, PALONOSETRON HYDROCHLORIDE
 PEMETREXED DISODIUM, PEMETREXED DISODIUM
 TADALAFIL, TADALAFIL

QINGDAO BAHEAL PHARM

* QINGDAO BAHEAL PHARMACEUTICAL CO LTD
 CELECOXIB, CELECOXIB
 DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE
 FOLIC ACID, FOLIC ACID
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 METHIMAZOLE, METHIMAZOLE

QOL MEDCL

* QOL MEDICAL LLC
 ETHAMOLIN, ETHANOLAMINE OLEATE

QUAGEN

* QUAGEN PHARMACEUTICALS LLC
 ALBUTEROL SULFATE, ALBUTEROL SULFATE
 BUTALBITAL AND ACETAMINOPHEN, ACETAMINOPHEN
 BUTALBITAL, ACETAMINOPHEN AND CAFFEINE, ACETAMINOPHEN
 CHILDREN'S CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
 CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
 CYPROHEPTADINE HYDROCHLORIDE, CYPROHEPTADINE HYDROCHLORIDE
 ESTRADIOL, ESTRADIOL
 FLUOCINOLONE ACETONIDE, FLUOCINOLONE ACETONIDE
 GLYCOPYRROLATE, GLYCOPYRROLATE
 HALOBETASOL PROPIONATE, HALOBETASOL PROPIONATE
 LEVETIRACETAM, LEVETIRACETAM
 LIDOCAINE, LIDOCAINE
 METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
 OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
 PROMETHAZINE HYDROCHLORIDE AND CODEINE PHOSPHATE, CODEINE PHOSPHATE
 PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
 PROPYLTHIOURACIL, PROPYLTHIOURACIL
 TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE
 VALPROIC ACID, VALPROIC ACID

QUEEN HAMAMATSU PET

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** Q ****

* QUEEN HAMAMATSU PET IMAGING CENTER
FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18

**** R ******R-PHARM US LLC**

* R-PHARM US LLC
IXEMPRA KIT, IXABEPILONE

RADIOMEDIX

* RADIOMEDIX INC
DETECTNET, COPPER CU-64 DOTATATE

RADIUS

* RADIUS HEALTH INC
BINOSTO, ALENDRONATE SODIUM
TYMLOS, ABALOPARATIDE

RAFA LABS LTD

* RAFA LABORATORIES LTD
MIDAZOLAM HYDROCHLORIDE (AUTOINJECTOR), MIDAZOLAM HYDROCHLORIDE

RAYNER SURGICAL

* RAYNER SURGICAL INC
OMIDRIA, KETOROLAC TROMETHAMINE

RB HLTH

* RB HEALTH US LLC
DELSYM, DEXTROMETHORPHAN POLISTIREX (OTC)
MUCINEX D, GUAIFENESIN (OTC)
MUCINEX DM, DEXTROMETHORPHAN HYDROBROMIDE (OTC)
MUCINEX, GUAIFENESIN (OTC)

REATA PHARMS

* REATA PHARMACEUTICALS INC
SKYCLARYS, OMAVELOXOLONE

RECORDATI RARE

* RECORDATI RARE DISEASES INC
CARBAGLU, CARGLUMIC ACID
CHEMET, SUCCIMER
CYSTADANE, BETAINE
CYSTADROPS, CYSTEAMINE HYDROCHLORIDE
ISTURISA, OSILODROSTAT PHOSPHATE
NEOPROFEN, IBUPROFEN LYSINE
SIGNIFOR LAR KIT, PASIREOTIDE PAMOATE
SIGNIFOR, PASIREOTIDE DIASPARTATE

REDHILL

* REDHILL BIOPHARMA LTD
TALICIA, AMOXICILLIN

RELIANCE LIFE

* RELIANCE LIFE SCIENCES PVT LTD
CAPECITABINE, CAPECITABINE
PEMETREXED DISODIUM, PEMETREXED DISODIUM

RELIANCE LIFE SCI

* RELIANCE LIFE SCIENCES PRIVATE LTD
BORTEZOMIB, BORTEZOMIB

REMPEX

* REMPEX PHARMACEUTICALS INC A WHOLLY OWNED SUB OF MELINTA THERAPEUTICS LLC
MINOCIN, MINOCYCLINE HYDROCHLORIDE
VABOMERE, MEROPENEM

RENATA

* RENATA LTD
FROVATRIPTAN SUCCINATE, FROVATRIPTAN SUCCINATE
MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
METOLAZONE, METOLAZONE
METOPROLOL TARTRATE, METOPROLOL TARTRATE
PREGABALIN, PREGABALIN
RISPERIDONE, RISPERIDONE
ROSUVASTATIN CALCIUM, ROSUVASTATIN CALCIUM

RENEW PHARMS

* RENEW PHARMACEUTICALS LTD

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** R ****

* RENEW PHARMACEUTICALS LTD
INDOCYANINE GREEN, INDOCYANINE GREEN

RESILIA PHARMS

* RESILIA PHARMACEUTICALS INC
ECOZA, ECONAZOLE NITRATE

REYOUNG

* REYOUNG CORPORATION
FENOFIBRATE (MICRONIZED), FENOFIBRATE
NEBIVOLOL HYDROCHLORIDE, NEBIVOLOL HYDROCHLORIDE
SERTRALINE HYDROCHLORIDE, SERTRALINE HYDROCHLORIDE
SILDENAFIL CITRATE, SILDENAFIL CITRATE
TENOFIVIR DISOPROXIL FUMARATE, TENOFIVIR DISOPROXIL FUMARATE

RHODES PHARMS

* RHODES PHARMACEUTICALS LP
AMPHETAMINE SULFATE, AMPHETAMINE SULFATE
APTENSIO XR, METHYLPHENIDATE HYDROCHLORIDE
BUPRENORPHINE HYDROCHLORIDE AND NALOXONE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE
BUPRENORPHINE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE
DEXMETHYLPHENIDATE HYDROCHLORIDE, DEXMETHYLPHENIDATE HYDROCHLORIDE
DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE
DILAUDID, HYDROMORPHONE HYDROCHLORIDE
FENOFIBRATE (MICRONIZED), FENOFIBRATE
FENOFIBRATE, FENOFIBRATE
HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
LISDEXAMFETAMINE DIMESYLATE, LISDEXAMFETAMINE DIMESYLATE
MORPHINE SULFATE, MORPHINE SULFATE
OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN
OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
SCOPOLAMINE, SCOPOLAMINE
THEOPHYLLINE, THEOPHYLLINE

RHYTHM

* RHYTHM PHARMACEUTICALS INC
IMCIVREE, SETMELANOTIDE ACETATE

RICONPHARMA LLC

* RICONPHARMA LLC
COLESTIPOL HYDROCHLORIDE, COLESTIPOL HYDROCHLORIDE
DICLOFENAC SODIUM, DICLOFENAC SODIUM
FLUNISOLIDE, FLUNISOLIDE
ISOSORBIDE DINITRATE AND HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
ISOSORBIDE MONONITRATE, ISOSORBIDE MONONITRATE
SCOPOLAMINE, SCOPOLAMINE

RIGEL PHARMS

* RIGEL PHARMACEUTICALS INC
GAVRETO, PRALSETINIB
REZLIDHIA, OLUTASIDENIB
TAVALISSE, FOSTAMATINIB DISODIUM

RILEY CONSUMER

* RILEY CONSUMER CARE LLC DBA CARLIN CONSUMER HEALTH
ZEGERID OTC, OMEPRAZOLE (OTC)

RISING

* RISING PHARMA HOLDING INC
AMILORIDE HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, AMILORIDE HYDROCHLORIDE
BUTORPHANOL TARTRATE, BUTORPHANOL TARTRATE
CAPECITABINE, CAPECITABINE
CAPTOPRIL AND HYDROCHLOROTHIAZIDE, CAPTOPRIL
CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE
CYTARABINE, CYTARABINE
DIGOXIN, DIGOXIN
DIPYRIDAMOLE, DIPYRIDAMOLE
DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
FLUMAZENIL, FLUMAZENIL
FLUNISOLIDE, FLUNISOLIDE
METHYLDOPA, METHYLDOPA
MYCOPHENOLATE MOFETIL, MYCOPHENOLATE MOFETIL
PENTOXIFYLLINE, PENTOXIFYLLINE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** R ****

* RISING PHARMA HOLDING INC
 PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE
 PYRIDOSTIGMINE BROMIDE, PYRIDOSTIGMINE BROMIDE
 TRAMADOL HYDROCHLORIDE AND ACETAMINOPHEN, ACETAMINOPHEN

* RISING PHARMA HOLDINGS INC
 ABIRATERONE ACETATE, ABIRATERONE ACETATE
 ACETIC ACID, ACETIC ACID, GLACIAL
 ACETYLCYSTEINE, ACETYLCYSTEINE
 ADENOSINE, ADENOSINE
 AKPENTOLATE, CYCLOPENTOLATE HYDROCHLORIDE
 ALBUTEROL SULFATE, ALBUTEROL SULFATE
 ALFENTA, ALFENTANIL HYDROCHLORIDE
 ALOSETRON HYDROCHLORIDE, ALOSETRON HYDROCHLORIDE
 AMANTADINE HYDROCHLORIDE, AMANTADINE HYDROCHLORIDE
 AMITRIPTYLINE HYDROCHLORIDE, AMITRIPTYLINE HYDROCHLORIDE
 APRACLONIDINE HYDROCHLORIDE, APRACLONIDINE HYDROCHLORIDE
 ATROPINE SULFATE, ATROPINE SULFATE
 AZATHIOPRINE, AZATHIOPRINE
 BACLOFEN, BACLOFEN
 BRIMONIDINE TARTRATE, BRIMONIDINE TARTRATE
 BUDESONIDE, BUDESONIDE
 BUMETANIDE, BUMETANIDE
 CALCITRIOL, CALCITRIOL
 CARBIDOPA, LEVODOPA AND ENTACAPONE, CARBIDOPA
 CEVIMELINE HYDROCHLORIDE, CEVIMELINE HYDROCHLORIDE
 CHLORZOXAZONE, CHLORZOXAZONE
 CICLOPIROX, CICLOPIROX
 CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE
 CLINDAMYCIN PALMITATE HYDROCHLORIDE, CLINDAMYCIN PALMITATE HYDROCHLORIDE
 CLINDAMYCIN PHOSPHATE IN 5% DEXTROSE IN PLASTIC CONTAINER, CLINDAMYCIN PHOSPHATE
 CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
 CLOPIDOGREL BISULFATE, CLOPIDOGREL BISULFATE
 CROMOLYN SODIUM, CROMOLYN SODIUM
 CYPROHEPTADINE HYDROCHLORIDE, CYPROHEPTADINE HYDROCHLORIDE
 DAPSONE, DAPSONE
 DESOXIMETASONE, DESOXIMETASONE
 DEXAMETHASONE, DEXAMETHASONE
 DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE
 DICLOFENAC SODIUM, DICLOFENAC SODIUM
 DICYCLOMINE HYDROCHLORIDE, DICYCLOMINE HYDROCHLORIDE
 DIFLORASONE DIACETATE, DIFLORASONE DIACETATE
 DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
 DOXAZOSIN MESYLATE, DOXAZOSIN MESYLATE
 DOXERCALCIFEROL, DOXERCALCIFEROL
 EPLERENONE, EPLERENONE
 ERLOTINIB HYDROCHLORIDE, ERLOTINIB HYDROCHLORIDE
 ERYTHROMYCIN AND BENZOYL PEROXIDE, BENZOYL PEROXIDE
 EXEMESTANE, EXEMESTANE
 FAMOTIDINE, FAMOTIDINE
 FENOFIBRATE, FENOFIBRATE
 FENOFIBRIC ACID, CHOLINE FENOFIBRATE
 FENOPROFEN CALCIUM, FENOPROFEN CALCIUM
 FEXOFENADINE HYDROCHLORIDE ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
 FEXOFENADINE HYDROCHLORIDE HIVES, FEXOFENADINE HYDROCHLORIDE (OTC)
 FEXOFENADINE HYDROCHLORIDE, FEXOFENADINE HYDROCHLORIDE
 FLUOCINOLONE ACETONIDE, FLUOCINOLONE ACETONIDE
 FLUPHENAZINE HYDROCHLORIDE, FLUPHENAZINE HYDROCHLORIDE
 FLURAZEPAM HYDROCHLORIDE, FLURAZEPAM HYDROCHLORIDE
 GABAPENTIN, GABAPENTIN
 GLYCOPYRROLATE, GLYCOPYRROLATE
 GRANISETRON HYDROCHLORIDE, GRANISETRON HYDROCHLORIDE
 HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
 HYDROCORTISONE, HYDROCORTISONE
 HYDROMORPHONE HYDROCHLORIDE, HYDROMORPHONE HYDROCHLORIDE
 INDAPAMIDE, INDAPAMIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** R ****

* RISING PHARMA HOLDINGS INC
 LABETALOL HYDROCHLORIDE, LABETALOL HYDROCHLORIDE
 LANSOPRAZOLE, AMOXICILLIN AND CLARITHROMYCIN (COPACKAGED), AMOXICILLIN
 LEVETIRACETAM, LEVETIRACETAM
 LEVOCARNITINE, LEVOCARNITINE
 LEVOFLOXACIN, LEVOFLOXACIN
 LINEZOLID, LINEZOLID
 LORAZEPAM, LORAZEPAM
 MESALAMINE, MESALAMINE
 METHIMAZOLE, METHIMAZOLE
 METHYLERGONOVINE MALEATE, METHYLERGONOVINE MALEATE
 METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
 MEXILETINE HYDROCHLORIDE, MEXILETINE HYDROCHLORIDE
 MONTELUKAST SODIUM, MONTELUKAST SODIUM
 MYCOPHENOLATE MOFETIL HYDROCHLORIDE, MYCOPHENOLATE MOFETIL HYDROCHLORIDE
 NADOLOL, NADOLOL
 NALOXONE HYDROCHLORIDE, NALOXONE HYDROCHLORIDE
 NAPHAZOLINE HYDROCHLORIDE AND PHENIRAMINE MALEATE, NAPHAZOLINE HYDROCHLORIDE (OTC)
 NATEGLINIDE, NATEGLINIDE
 NEMBUTAL SODIUM, PENTOBARBITAL SODIUM
 NITAZOXANIDE, NITAZOXANIDE
 NOREPINEPHRINE BITARTRATE, NOREPINEPHRINE BITARTRATE
 NYSTATIN AND TRIAMCINOLONE ACETONIDE, NYSTATIN
 ORPHENADRINE CITRATE, ORPHENADRINE CITRATE
 OXYBUTYNYN CHLORIDE, OXYBUTYNYN CHLORIDE
 PARICALCITOL, PARICALCITOL
 PERPHENAZINE, PERPHENAZINE
 PHENOXYBENZAMINE HYDROCHLORIDE, PHENOXYBENZAMINE HYDROCHLORIDE
 PHENYTOIN, PHENYTOIN
 PILOCARPINE HYDROCHLORIDE, PILOCARPINE HYDROCHLORIDE
 POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
 POTASSIUM CITRATE, POTASSIUM CITRATE
 PREGABALIN, PREGABALIN
 PROBENECID AND COLCHICINE, COLCHICINE
 PROBENECID, PROBENECID
 PROPAFENONE HYDROCHLORIDE, PROPAFENONE HYDROCHLORIDE
 PROPARACAINE HYDROCHLORIDE, PROPARACAINE HYDROCHLORIDE
 PYRIDOSTIGMINE BROMIDE, PYRIDOSTIGMINE BROMIDE
 RANOLAZINE, RANOLAZINE
 RISPERIDONE, RISPERIDONE
 ROPIVACAINE HYDROCHLORIDE, ROPIVACAINE HYDROCHLORIDE
 SELEGILINE HYDROCHLORIDE, SELEGILINE HYDROCHLORIDE
 SEVELAMER CARBONATE, SEVELAMER CARBONATE
 SOLIFENACIN SUCCINATE, SOLIFENACIN SUCCINATE
 SUBLIMAZE PRESERVATIVE FREE, FENTANYL CITRATE
 SUFENTA PRESERVATIVE FREE, SUFENTANIL CITRATE
 SULFAMYLON, MAFENIDE ACETATE
 SUMATRIPTAN AND NAPROXEN SODIUM, NAPROXEN SODIUM
 TEMOZOLOMIDE, TEMOZOLOMIDE
 THIOTHIXENE, THIOTHIXENE
 TIMOLOL MALEATE, TIMOLOL MALEATE
 TOLMETIN SODIUM, TOLMETIN SODIUM
 TOREMIFENE CITRATE, TOREMIFENE CITRATE
 TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE
 TRIENTINE HYDROCHLORIDE, TRIENTINE HYDROCHLORIDE
 TROPICACYL, TROPICAMIDE
 VORICONAZOLE, VORICONAZOLE
 ZILEUTON, ZILEUTON

* RISING PHARMACEUTICALS
 URSODIOL, URSODIOL

RISING PHARMS

* RISING PHARMACEUTICALS INC
 ZAFIRLUKAST, ZAFIRLUKAST

RK PHARMA

* RK PHARMA INC

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** R ******* RK PHARMA INC**

CHROMIC CHLORIDE, CHROMIC CHLORIDE
 CUPRIC CHLORIDE, CUPRIC CHLORIDE
 DICLOFENAC POTASSIUM, DICLOFENAC POTASSIUM
 DOXEPIN HYDROCHLORIDE, DOXEPIN HYDROCHLORIDE
 LEUPROLIDE ACETATE, LEUPROLIDE ACETATE
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 METHYLENE BLUE, METHYLENE BLUE
 MITOMYCIN, MITOMYCIN
 MYCOPHENOLIC SODIUM, MYCOPHENOLIC SODIUM
 NADOLOL, NADOLOL
 NICARDIPINE HYDROCHLORIDE, NICARDIPINE HYDROCHLORIDE
 PALIPERIDONE, PALIPERIDONE
 URSODIOL, URSODIOL

ROCHE PALO*** ROCHE PALO ALTO LLC**

CELLCEPT, MYCOPHENOLATE MOFETIL
 CELLCEPT, MYCOPHENOLATE MOFETIL HYDROCHLORIDE

RUBICON*** RUBICON RESEARCH PRIVATE LTD**

ACETAZOLAMIDE, ACETAZOLAMIDE
 ACYCLOVIR, ACYCLOVIR
 AMANTADINE HYDROCHLORIDE, AMANTADINE HYDROCHLORIDE
 AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE
 AMITRIPTYLINE HYDROCHLORIDE, AMITRIPTYLINE HYDROCHLORIDE
 ARIPIPIRAZOLE, ARIPIPIRAZOLE
 BACLOFEN, BACLOFEN
 BISOPROLOL FUMARATE, BISOPROLOL FUMARATE
 BUMETANIDE, BUMETANIDE
 BUPRENORPHINE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE
 BUSPIRONE HYDROCHLORIDE, BUSPIRONE HYDROCHLORIDE
 CARBIDOPA AND LEVODOPA, CARBIDOPA
 CARVEDILOL, CARVEDILOL
 CEVIMELINE HYDROCHLORIDE, CEVIMELINE HYDROCHLORIDE
 CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE
 CLOMIPHENE CITRATE, CLOMIPHENE CITRATE
 CLONAZEPAM, CLONAZEPAM
 CYCLOBENZAPRINE HYDROCHLORIDE, CYCLOBENZAPRINE HYDROCHLORIDE
 DESVENLAFAXINE SUCCINATE, DESVENLAFAXINE SUCCINATE
 DICLOFENAC POTASSIUM, DICLOFENAC POTASSIUM
 DICLOFENAC SODIUM, DICLOFENAC SODIUM
 DICYCLOMINE HYDROCHLORIDE, DICYCLOMINE HYDROCHLORIDE
 DIHYDROERGOTAMINE MESYLATE, DIHYDROERGOTAMINE MESYLATE
 DORZOLAMIDE HYDROCHLORIDE, DORZOLAMIDE HYDROCHLORIDE
 GABAPENTIN, GABAPENTIN
 GLIPIZIDE, GLIPIZIDE
 GUANFACINE HYDROCHLORIDE, GUANFACINE HYDROCHLORIDE
 HALOPERIDOL, HALOPERIDOL LACTATE
 IPRATROPIUM BROMIDE, IPRATROPIUM BROMIDE
 ISOSORBIDE DINITRATE, ISOSORBIDE DINITRATE
 LABETALOL HYDROCHLORIDE, LABETALOL HYDROCHLORIDE
 LAMOTRIGINE, LAMOTRIGINE
 LEVOFLOXACIN, LEVOFLOXACIN
 LIDOCAINE HYDROCHLORIDE, LIDOCAINE HYDROCHLORIDE
 LITHIUM CITRATE, LITHIUM CITRATE
 LOPERAMIDE HYDROCHLORIDE, LOPERAMIDE HYDROCHLORIDE
 LORATADINE, LORATADINE (OTC)
 METOLAZONE, METOLAZONE
 METOPROLOL TARTRATE, METOPROLOL TARTRATE
 MIDODRINE HYDROCHLORIDE, MIDODRINE HYDROCHLORIDE
 NITROGLYCERIN, NITROGLYCERIN
 NORTRIPTYLINE HYDROCHLORIDE, NORTRIPTYLINE HYDROCHLORIDE
 OXCARBAZEPINE, OXCARBAZEPINE
 OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE
 PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** R ****

* RUBICON RESEARCH PRIVATE LTD
 POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
 PREGABALIN, PREGABALIN
 PRIMIDONE, PRIMIDONE
 RABEPRAZOLE SODIUM, RABEPRAZOLE SODIUM
 SILDENAFIL CITRATE, SILDENAFIL CITRATE
 TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE
 TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE
 TRANEXAMIC ACID, TRANEXAMIC ACID
 TRIAMTERENE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE

RVL PHARMS

* RVL PHARMACEUTICALS INC
 UPNEEQ, OXYMETAZOLINE HYDROCHLORIDE

RYAN LABS

* RYAN LABORATORIES LLC
 VALSARTAN, VALSARTAN
 VIGABATRIN, VIGABATRIN

**** S ******SABA ILAC SANAYIVE**

* SABA ILAC SANAYIVE TICARET AS
 FUROSEMIDE, FUROSEMIDE

SAGE PRODS

* SAGE PRODUCTS INC
 CHLORHEXIDINE GLUCONATE, CHLORHEXIDINE GLUCONATE (OTC)

SAGE THERAP

* SAGE THERAPEUTICS INC
 ZULRESSO, BREXANOLONE

SAGENT

* SAGENT PHARMACEUTICALS
 BUMETANIDE, BUMETANIDE
 DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
 FAMOTIDINE PRESERVATIVE FREE, FAMOTIDINE
 FAMOTIDINE, FAMOTIDINE
 FUROSEMIDE, FUROSEMIDE
 GLYCOPYRROLATE, GLYCOPYRROLATE
 PROCHLORPERAZINE EDISYLATE, PROCHLORPERAZINE EDISYLATE
 VALPROATE SODIUM, VALPROATE SODIUM

SAGENT PHARMS

* SAGENT PHARMACEUTICALS INC
 CAFFEINE CITRATE, CAFFEINE CITRATE
 DOXORUBICIN HYDROCHLORIDE, DOXORUBICIN HYDROCHLORIDE
 FLUMAZENIL, FLUMAZENIL
 HALOPERIDOL, HALOPERIDOL LACTATE
 HEPARIN SODIUM PRESERVATIVE FREE, HEPARIN SODIUM
 HEPARIN SODIUM, HEPARIN SODIUM
 LEUCOVORIN CALCIUM PRESERVATIVE FREE, LEUCOVORIN CALCIUM
 LEVETIRACETAM, LEVETIRACETAM
 NAFCILLIN SODIUM, NAFCILLIN SODIUM
 ORPHENADRINE CITRATE, ORPHENADRINE CITRATE
 OXACILLIN SODIUM, OXACILLIN SODIUM
 TOPOTECAN HYDROCHLORIDE, TOPOTECAN HYDROCHLORIDE
 VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE

SAGENT PHARMS INC

* SAGENT PHARMACEUTICALS INC
 ACETYLCYSTEINE, ACETYLCYSTEINE
 AMIKACIN SULFATE, AMIKACIN SULFATE
 AMPICILLIN SODIUM, AMPICILLIN SODIUM
 CEFEPIME HYDROCHLORIDE, CEFEPIME HYDROCHLORIDE
 CHLOROTHIAZIDE SODIUM, CHLOROTHIAZIDE SODIUM
 CISATRACURIUM BESYLATE, CISATRACURIUM BESYLATE
 CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
 COLISTIMETHATE SODIUM, COLISTIMETHATE SODIUM
 CYANOCOBALAMIN, CYANOCOBALAMIN
 CYCLOPHOSPHAMIDE, CYCLOPHOSPHAMIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** S ****

* SAGENT PHARMACEUTICALS INC
 DAPTOMYCIN, DAPTOMYCIN
 DECITABINE, DECITABINE
 DESMOPRESSIN ACETATE, DESMOPRESSIN ACETATE
 DIHYDROERGOTAMINE MESYLATE, DIHYDROERGOTAMINE MESYLATE
 EPHEDRINE SULFATE, EPHEDRINE SULFATE
 EPTIFIBATIDE, EPTIFIBATIDE
 ESMOLOL HYDROCHLORIDE, ESMOLOL HYDROCHLORIDE
 FLUDARABINE PHOSPHATE, FLUDARABINE PHOSPHATE
 FLUOROURACIL, FLUOROURACIL
 FULVESTRANT, FULVESTRANT
 GEMCITABINE HYDROCHLORIDE, GEMCITABINE HYDROCHLORIDE
 GLYDO, LIDOCAINE HYDROCHLORIDE
 KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
 LEUCOVORIN CALCIUM PRESERVATIVE FREE, LEUCOVORIN CALCIUM
 LINEZOLID, LINEZOLID
 MELPHALAN HYDROCHLORIDE, MELPHALAN HYDROCHLORIDE
 MESNA, MESNA
 METHOCARBAMOL, METHOCARBAMOL
 METHOTREXATE SODIUM PRESERVATIVE FREE, METHOTREXATE SODIUM
 METHYLPREDNISOLONE ACETATE, METHYLPREDNISOLONE ACETATE
 METHYLPREDNISOLONE SODIUM SUCCINATE, METHYLPREDNISOLONE SODIUM SUCCINATE
 NEOSTIGMINE METHYLSULFATE, NEOSTIGMINE METHYLSULFATE
 OCTREOTIDE ACETATE (PRESERVATIVE FREE), OCTREOTIDE ACETATE
 OCTREOTIDE ACETATE, OCTREOTIDE ACETATE
 OXYTOCIN, OXYTOCIN
 PALONOSETRON HYDROCHLORIDE, PALONOSETRON HYDROCHLORIDE
 PAMIDRONATE DISODIUM, PAMIDRONATE DISODIUM
 PENTOBARBITAL SODIUM, PENTOBARBITAL SODIUM
 PHENYLEPHRINE HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE
 PIPERACILLIN AND TAZOBACTAM, PIPERACILLIN SODIUM
 PROPOFOL, PROPOFOL
 ROCURONIUM BROMIDE, ROCURONIUM BROMIDE
 SODIUM NITROPRUSSIDE, SODIUM NITROPRUSSIDE
 SUCCINYLCHOLINE CHLORIDE, SUCCINYLCHOLINE CHLORIDE
 THIAMINE HYDROCHLORIDE, THIAMINE HYDROCHLORIDE
 VECURONIUM BROMIDE, VECURONIUM BROMIDE
 ZOLEDRONIC ACID, ZOLEDRONIC ACID

SALERNO PHARMS

* SALERNO PHARMACEUTICALS
 FLOLIPID, SIMVASTATIN

SALIX

* SALIX PHARMACEUTICALS INC
 APRISO, MESALAMINE
 PLENVU, ASCORBIC ACID
 RELISTOR, METHYLNALTREXONE BROMIDE
 TRULANCE, PLECANATIDE
 UCERIS, BUDESONIDE

SALIX PHARMS

* SALIX PHARMACEUTICALS INC
 ANUSOL HC, HYDROCORTISONE
 DIURIL, CHLOROTHIAZIDE
 MOVIPREP, ASCORBIC ACID
 RELISTOR, METHYLNALTREXONE BROMIDE
 XIFAXAN, RIFAXIMIN

SAMSON MEDCL

* SAMSON MEDICAL TECHNOLOGIES LLC
 CEFAZOLIN SODIUM, CEFAZOLIN SODIUM
 CEFEPIME HYDROCHLORIDE IN PLASTIC CONTAINER, CEFEPIME HYDROCHLORIDE
 CEFOXITIN IN PLASTIC CONTAINER, CEFOXITIN SODIUM
 CEFTRIAZONE, CEFTRIAZONE SODIUM
 VANCOMYCIN HYDROCHLORIDE IN PLASTIC CONTAINER, VANCOMYCIN HYDROCHLORIDE

SANALUZ

* SANALUZ LLC
 AMPHETAMINE SULFATE, AMPHETAMINE SULFATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** S ******* SANALUZ LLC**

PYRIMETHAMINE, PYRIMETHAMINE
SEROMYCIN, CYCLOSERINE

SANDOZ*** SANDOZ**

DOCETAXEL, DOCETAXEL

*** SANDOZ INC**

ACETAMINOPHEN, ACETAMINOPHEN
ALBUTEROL SULFATE, ALBUTEROL SULFATE
ALPRAZOLAM, ALPRAZOLAM
AMANTADINE HYDROCHLORIDE, AMANTADINE HYDROCHLORIDE
AMITRIPTYLINE HYDROCHLORIDE, AMITRIPTYLINE HYDROCHLORIDE
AMOXICILLIN AND CLAVULANATE POTASSIUM, AMOXICILLIN
AMOXICILLIN, AMOXICILLIN
AMPICILLIN AND SULBACTAM, AMPICILLIN SODIUM
AMPICILLIN SODIUM, AMPICILLIN SODIUM
AMPICILLIN TRIHYDRATE, AMPICILLIN/AMPICILLIN TRIHYDRATE
ANECTINE, SUCCINYLCHOLINE CHLORIDE
ANGIOMAX, BIVALIRUDIN
APREPITANT, APREPITANT
ARGATROBAN IN SODIUM CHLORIDE, ARGATROBAN
ARRANON, NELARABINE
ARSENIC TRIOXIDE, ARSENIC TRIOXIDE
ASPIRIN AND DIPYRIDAMOLE, ASPIRIN
AZELASTINE HYDROCHLORIDE, AZELASTINE HYDROCHLORIDE
AZITHROMYCIN, AZITHROMYCIN
AZOPT, BRINZOLAMIDE
BENZAEPRILOL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, BENAZEPRIL HYDROCHLORIDE
BETOPTIC, BETAXOLOL HYDROCHLORIDE
BICALUTAMIDE, BICALUTAMIDE
BIMATOPROST, BIMATOPROST
BORTEZOMIB, BORTEZOMIB
BRIMONIDINE TARTRATE AND TIMOLOL MALEATE, BRIMONIDINE TARTRATE
BRIMONIDINE TARTRATE, BRIMONIDINE TARTRATE
BROMOCRIPTINE MESYLATE, BROMOCRIPTINE MESYLATE
BUDESONIDE, BUDESONIDE
BUMETANIDE, BUMETANIDE
BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
CARBOPLATIN, CARBOPLATIN
CARTEOLOL HYDROCHLORIDE, CARTEOLOL HYDROCHLORIDE
CARVEDILOL, CARVEDILOL
CEFAZOLIN SODIUM, CEFZOLIN SODIUM
CEFPODOXIME PROXETIL, CEFPODOXIME PROXETIL
CEFTRIAZONE, CEFTRIAZONE SODIUM
CHILDREN'S CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
CILLOXAN, CIPROFLOXACIN HYDROCHLORIDE
CIPRO HC, CIPROFLOXACIN HYDROCHLORIDE
CIPRODEX, CIPROFLOXACIN
CLARITHROMYCIN, CLARITHROMYCIN
CLOMIPRAMINE HYDROCHLORIDE, CLOMIPRAMINE HYDROCHLORIDE
CLONAZEPAM, CLONAZEPAM
COSYNTROPIN, COSYNTROPIN
CROMOLYN SODIUM, CROMOLYN SODIUM
CYANOCOBALAMIN, CYANOCOBALAMIN
CYCLOPHOSPHAMIDE, CYCLOPHOSPHAMIDE
CYCLOSPORINE, CYCLOSPORINE
DECITABINE, DECITABINE
DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE
DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE
DICLOFENAC SODIUM, DICLOFENAC SODIUM
DIGOXIN, DIGOXIN
DORZOLAMIDE HYDROCHLORIDE AND TIMOLOL MALEATE, DORZOLAMIDE HYDROCHLORIDE
DORZOLAMIDE HYDROCHLORIDE, DORZOLAMIDE HYDROCHLORIDE
DOXERCALCIFEROL, DOXERCALCIFEROL
DUREZOL, DIFLUPREDNATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** S ****

* SANDOZ INC
 EDARAVONE, EDARAVONE
 ENOXAPARIN SODIUM (PRESERVATIVE FREE), ENOXAPARIN SODIUM
 EPHEDRINE SULFATE, EPHEDRINE SULFATE
 EPLERENONE, EPLERENONE
 EXELON, RIVASTIGMINE
 EZETIMIBE, EZETIMIBE
 FERUMOXYTOL, FERUMOXYTOL
 FLUMAZENIL, FLUMAZENIL
 FOCALIN XR, DEXMETHYLPHENIDATE HYDROCHLORIDE
 FOCALIN, DEXMETHYLPHENIDATE HYDROCHLORIDE
 FOSINOPRIL SODIUM AND HYDROCHLOROTHIAZIDE, FOSINOPRIL SODIUM
 FULVESTRANT, FULVESTRANT
 GALANTAMINE HYDROBROMIDE, GALANTAMINE HYDROBROMIDE
 GATIFLOXACIN, GATIFLOXACIN
 GENTAMICIN SULFATE, GENTAMICIN SULFATE
 GLATOPA, GLATIRAMER ACETATE
 GLYCOPYRROLATE, GLYCOPYRROLATE
 GRANISETRON HYDROCHLORIDE, GRANISETRON HYDROCHLORIDE
 GRISEOFULVIN, GRISEOFULVIN, MICROSIZE
 GRISEOFULVIN, ULTRAMICROSIZE, GRISEOFULVIN, ULTRAMICROSIZE
 GUANFACINE HYDROCHLORIDE, GUANFACINE HYDROCHLORIDE
 HEPARIN SODIUM, HEPARIN SODIUM
 HYCAMTIN, TOPOTECAN HYDROCHLORIDE
 HYDROXYCHLOROQUINE SULFATE, HYDROXYCHLOROQUINE SULFATE
 HYDROXYZINE PAMOATE, HYDROXYZINE PAMOATE
 IMIPRAMINE HYDROCHLORIDE, IMIPRAMINE HYDROCHLORIDE
 INDOMETHACIN, INDOMETHACIN
 IRBESARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 IRBESARTAN, IRBESARTAN
 ISONIAZID, ISONIAZID
 ISOPTO CARPINE, PILOCARPINE HYDROCHLORIDE
 ISOSORBIDE DINITRATE, ISOSORBIDE DINITRATE
 ITRACONAZOLE, ITRACONAZOLE
 KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
 LABETALOL HYDROCHLORIDE, LABETALOL HYDROCHLORIDE
 LANORINAL, ASPIRIN
 LANSOPRAZOLE, AMOXICILLIN AND CLARITHROMYCIN (COPACKAGED), AMOXICILLIN
 LATANOPROST, LATANOPROST
 LESCOL XL, FLUVASTATIN SODIUM
 LEUPROLIDE ACETATE, LEUPROLIDE ACETATE
 LEVOLEUCOVORIN CALCIUM, LEVOLEUCOVORIN CALCIUM
 LINEZOLID, LINEZOLID
 LISDEXAMFETAMINE DIMESYLATE, LISDEXAMFETAMINE DIMESYLATE
 LISINAPRIL AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 LORAZEPAM, LORAZEPAM
 LOTREL, AMLODIPINE BESYLATE
 MAXITROL, DEXAMETHASONE
 MECLIZINE HYDROCHLORIDE, MECLIZINE HYDROCHLORIDE
 MESALAMINE, MESALAMINE
 METAXALONE, METAXALONE
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 METHOTREXATE SODIUM PRESERVATIVE FREE, METHOTREXATE SODIUM
 METHYLPREDNISOLONE ACETATE, METHYLPREDNISOLONE ACETATE
 METHYLPREDNISOLONE, METHYLPREDNISOLONE
 METOLAZONE, METOLAZONE
 METOPROLOL TARTRATE, METOPROLOL TARTRATE
 MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE
 MYCOPHENOLATE MOFETIL, MYCOPHENOLATE MOFETIL
 MYDRIACYL, TROPICAMIDE
 NADOLOL, NADOLOL
 NEOMYCIN AND POLYMYXIN B SULFATES AND HYDROCORTISONE, HYDROCORTISONE
 NEVIRAPINE, NEVIRAPINE
 NITROFURANTOIN (MONOHYDRATE/MACROCRYSTALS), NITROFURANTOIN
 NOREPINEPHRINE BITARTRATE, NOREPINEPHRINE BITARTRATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** S ****

* SANDOZ INC
 OMEPRAZOLE, OMEPRAZOLE
 OMNIPRED, PREDNISOLONE ACETATE
 ONDANSETRON HYDROCHLORIDE PRESERVATIVE FREE, ONDANSETRON HYDROCHLORIDE
 ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
 ONDANSETRON, ONDANSETRON
 ORPHENADRINE CITRATE, ORPHENADRINE CITRATE
 OXALIPLATIN, OXALIPLATIN
 PALONOSETRON HYDROCHLORIDE, PALONOSETRON HYDROCHLORIDE
 PARICALCITOL, PARICALCITOL
 PEMETREXED DISODIUM, PEMETREXED DISODIUM
 PENICILLIN G POTASSIUM, PENICILLIN G POTASSIUM
 PENICILLIN G SODIUM, PENICILLIN G SODIUM
 PERPHENAZINE, PERPHENAZINE
 PHENTERMINE HYDROCHLORIDE, PHENTERMINE HYDROCHLORIDE
 PHENYLEPHRINE HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE
 PIOGLITAZONE HYDROCHLORIDE, PIOGLITAZONE HYDROCHLORIDE
 PIPERACILLIN AND TAZOBACTAM, PIPERACILLIN SODIUM
 PIRFENIDONE, PIRFENIDONE
 PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
 QOLIANA, BRIMONIDINE TARTRATE
 RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
 RECLAST, ZOLEDRONIC ACID
 REGONOL, PYRIDOSTIGMINE BROMIDE
 RIBAVIRIN, RIBAVIRIN
 RISPERIDONE, RISPERIDONE
 RITALIN LA, METHYLPHENIDATE HYDROCHLORIDE
 RITALIN, METHYLPHENIDATE HYDROCHLORIDE
 ROCURONIUM BROMIDE, ROCURONIUM BROMIDE
 TACROLIMUS, TACROLIMUS
 TAFLUPROST, TAFLUPROST
 TAMSULOSIN HYDROCHLORIDE, TAMSULOSIN HYDROCHLORIDE
 TELMISARTAN, TELMISARTAN
 TERIFLUNOMIDE, TERIFLUNOMIDE
 TESTOSTERONE CYPIONATE, TESTOSTERONE CYPIONATE
 TIGECYCLINE, TIGECYCLINE
 TIMOLOL MALEATE, TIMOLOL MALEATE
 TOBRADEX, DEXAMETHASONE
 TOBREX, TOBRAMYCIN
 TRAVATAN Z, TRAVOPROST
 TREPROSTINIL, TREPROSTINIL
 TRIAMTERENE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 TRIFLUOPERAZINE HYDROCHLORIDE, TRIFLUOPERAZINE HYDROCHLORIDE
 TRIFLURIDINE, TRIFLURIDINE
 TRIMETHOPRIM SULFATE AND POLYMYXIN B SULFATE, POLYMYXIN B SULFATE
 VIVELLE-DOT, ESTRADIOL
 ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE

SANDOZ CANADA INC

* SANDOZ CANADA INC
 INFUVITE ADULT, ALPHA-TOCOPHEROL ACETATE
 INFUVITE PEDIATRIC (PHARMACY BULK PACKAGE), ASCORBIC ACID
 INFUVITE PEDIATRIC, ASCORBIC ACID

SANDOZ INC

* SANDOZ INC
 AMOXICILLIN AND CLAVULANATE POTASSIUM, AMOXICILLIN
 CEFTRIAZONE, CEFTRIAZONE SODIUM
 CISATRACURIUM BESYLATE PRESERVATIVE FREE, CISATRACURIUM BESYLATE
 CISATRACURIUM BESYLATE, CISATRACURIUM BESYLATE
 CLINDAMYCIN PHOSPHATE IN 5% DEXTROSE IN PLASTIC CONTAINER, CLINDAMYCIN PHOSPHATE
 ENALAPRIL MALEATE, ENALAPRIL MALEATE
 ENOXAPARIN SODIUM, ENOXAPARIN SODIUM
 GRANISETRON HYDROCHLORIDE, GRANISETRON HYDROCHLORIDE
 OLANZAPINE, OLANZAPINE
 ROPINIROLE HYDROCHLORIDE, ROPINIROLE HYDROCHLORIDE
 VORICONAZOLE, VORICONAZOLE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** S ****

* SANDOZ INC
ZIPRASIDONE HYDROCHLORIDE, ZIPRASIDONE HYDROCHLORIDE

SANOFI

* SANOFI AVENTIS US LLC
FEXINIDAZOLE, FEXINIDAZOLE
FLOMAX, TAMSULOSIN HYDROCHLORIDE

* SANOFI GENZYME
HECTOROL, DOXERCALCIFEROL
REVELA, SEVELAMER CARBONATE

SANOFI AVENTIS US

* SANOFI AVENTIS US INC
JEVTANA KIT, CABAZITAXEL

* SANOFI AVENTIS US LLC
AMARYL, GLIMEPIRIDE
AMBIEN CR, ZOLPIDEM TARTRATE
AMBIEN, ZOLPIDEM TARTRATE
ARAVA, LEFLUNOMIDE
AUBAGIO, TERIFLUNOMIDE
AVALIDE, HYDROCHLOROTHIAZIDE
AVAPRO, IRBESARTAN
DIABETA, GLYBURIDE
FERRLECIT, FERRIC OXYHYDROXIDE
LOVENOX (PRESERVATIVE FREE), ENOXAPARIN SODIUM
LOVENOX, ENOXAPARIN SODIUM
MULTAQ, DRONEDARONE HYDROCHLORIDE
PLAVIX, CLOPIDOGREL BISULFATE
PRIFTIN, RIFAPENTINE
PRIMAQUINE, PRIMAQUINE PHOSPHATE
RIFADIN, RIFAMPIN
TAXOTERE, DOCETAXEL

SANTARUS INC

* SANTARUS INC
GLUMETZA, METFORMIN HYDROCHLORIDE

SAPTALIS PHARMS

* SAPTALIS PHARMACEUTICALS LLC
CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
CYCLOSPORINE, CYCLOSPORINE
GLYCOPYRROLATE, GLYCOPYRROLATE
LIKMEZ, METRONIDAZOLE
METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
METRONIDAZOLE, METRONIDAZOLE
VOSOL HC, ACETIC ACID, GLACIAL

SAREPTA THERAPS INC

* SAREPTA THERAPEUTICS INC
AMONDYS 45, CASIMERSSEN
EXONDYS 51, ETEPLIRSEN
VYONDYS 53, GOLODIRSEN

SARFE PHARMS

* SARFEZ PHARMACEUTICALS INC
SOANZ, TORSEMIDE

SAVIOR LIFETEC CORP

* SAVIOR LIFETEC CORP
ERTAPENEM SODIUM, ERTAPENEM SODIUM
MEROPENEM, MEROPENEM

SAWAI USA

* SAWAI USA INC
PITAVASTATIN CALCIUM, PITAVASTATIN CALCIUM

SCHERING

* SCHERING CORP
NOXAFIL, POSACONAZOLE

SCIARRA LABS

* SCIARRA LABORATORIES INC
SCLEROSOL, TALC
TALC, TALC

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** S ******SCIECURE**

* SCIECURE LABORATORIES INC
FOSCARNET SODIUM, FOSCARNET SODIUM

SCIECURE PHARMA INC

* SCIECURE PHARMA INC
PAROXETINE HYDROCHLORIDE, PAROXETINE HYDROCHLORIDE

SCIEGEN PHARMS INC

* SCIEGEN PHARMACEUTICALS INC
ACETIC ACID, ACETIC ACID, GLACIAL
AMLODIPINE AND OLMESARTAN MEDOXOMIL, AMLODIPINE BESYLATE
ARIPIPIRAZOLE, ARIPIPIRAZOLE
ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM
BETAMETHASONE VALERATE, BETAMETHASONE VALERATE
BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
CARBIDOPA AND LEVODOPA, CARBIDOPA
CARISOPRODOL, CARISOPRODOL
CELECOXIB, CELECOXIB
CLOPIDOGREL BISULFATE, CLOPIDOGREL BISULFATE
CLOTRIMAZOLE, CLOTRIMAZOLE
DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
DROXIDOPA, DROXIDOPA
ETHACRYNIC ACID, ETHACRYNIC ACID
EZETIMIBE AND SIMVASTATIN, EZETIMIBE
EZETIMIBE, EZETIMIBE
FEXOFENADINE HYDROCHLORIDE ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
FEXOFENADINE HYDROCHLORIDE HIVES, FEXOFENADINE HYDROCHLORIDE (OTC)
FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
GABAPENTIN, GABAPENTIN
HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
IRBESARTAN, IRBESARTAN
LACOSAMIDE, LACOSAMIDE
LAMOTRIGINE, LAMOTRIGINE
LEVETIRACETAM, LEVETIRACETAM
LEVOCARNITINE, LEVOCARNITINE
LEVOCETIRIZINE DIHYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE
LISINOPRIL, LISINOPRIL
LITHIUM CITRATE, LITHIUM CITRATE
METAXALONE, METAXALONE
METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
NABUMETONE, NABUMETONE
NAPROXEN AND ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM
NAPROXEN SODIUM, NAPROXEN SODIUM
NAPROXEN, NAPROXEN
OLMESARTAN MEDOXOMIL, OLMESARTAN MEDOXOMIL
OMEPRAZOLE AND SODIUM BICARBONATE, OMEPRAZOLE
PHYTONADIONE, PHYTONADIONE
PIRFENIDONE, PIRFENIDONE
POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE
PREGABALIN, PREGABALIN
QUETIAPINE FUMARATE, QUETIAPINE FUMARATE
RALOXIFENE HYDROCHLORIDE, RALOXIFENE HYDROCHLORIDE
RANOLAZINE, RANOLAZINE
ROSUVASTATIN CALCIUM, ROSUVASTATIN CALCIUM
SERTRALINE HYDROCHLORIDE, SERTRALINE HYDROCHLORIDE
SOLIFENACIN SUCCINATE, SOLIFENACIN SUCCINATE
TAMSULOSIN HYDROCHLORIDE, TAMSULOSIN HYDROCHLORIDE
TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE
VALSARTAN, VALSARTAN

SCILEX PHARMS

* SCILEX PHARMACEUTICALS INC
ELYXYB, CELECOXIB
GLOPERBA, COLCHICINE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** S ****

* SCILEX PHARMACEUTICALS INC
ZTLIDO, LIDOCAINE

SCINOPHARM TAIWAN

* SCINOPHARM TAIWAN LTD
CLOFARABINE, CLOFARABINE
FONDAPARINUX SODIUM, FONDAPARINUX SODIUM

SCPHARMACEUTICALS

* SCPHARMACEUTICALS INC
FUROSCIX, FUROSEMIDE

SCYNEXIS

* SCYNEXIS INC
BREXAFEMME, IBREXAFUNGERP CITRATE

SEAGEN

* SEAGEN INC
TUKYSA, TUCATINIB

SECAN PHARMS

* SECAN PHARMACEUTICALS INC
LEVETIRACETAM, LEVETIRACETAM

SECURA

* SECURA BIO INC
COPIKTRA, DUVELISIB

SENORES PHARMS

* SENORES PHARMACEUTICALS INC
AMPHETAMINE SULFATE, AMPHETAMINE SULFATE
BUTALBITAL AND ACETAMINOPHEN, ACETAMINOPHEN
BUTALBITAL, ACETAMINOPHEN AND CAFFEINE, ACETAMINOPHEN
CHLORZOXAZONE, CHLORZOXAZONE
DICLOFENAC POTASSIUM, DICLOFENAC POTASSIUM
HYDROXYCHLOROQUINE SULFATE, HYDROXYCHLOROQUINE SULFATE
IVERMECTIN, IVERMECTIN
KETOCONAZOLE, KETOCONAZOLE
KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
MEXILETINE HYDROCHLORIDE, MEXILETINE HYDROCHLORIDE
NICARDIPINE HYDROCHLORIDE, NICARDIPINE HYDROCHLORIDE
TOPIRAMATE, TOPIRAMATE

SENTISS

* SENTISS AG
ALBUTEROL SULFATE, ALBUTEROL SULFATE
BRIMONIDINE TARTRATE AND TIMOLOL MALEATE, BRIMONIDINE TARTRATE
BROMFENAC SODIUM, BROMFENAC SODIUM
CIPROFLOXACIN AND DEXAMETHASONE, CIPROFLOXACIN
ERYTHROMYCIN, ERYTHROMYCIN
KETOTIFEN FUMARATE, KETOTIFEN FUMARATE (OTC)
LIDOCAINE HYDROCHLORIDE, LIDOCAINE HYDROCHLORIDE
LOTEPREDNOL ETABONATE, LOTEPREDNOL ETABONATE
OFLOXACIN, OFLOXACIN

SENTISS PHARMA

* SENTISS PHARMA PRIVATE LTD
TIMOLOL MALEATE, TIMOLOL MALEATE

SENTYNL THERAPS INC

* SENTYNL THERAPEUTICS INC
NULIBRY, FOSDENOPTERIN HYDROBROMIDE
ZOKINVY, LONAFARNIB

SEPTODONT

* SEPTODONT INC
DYCLOPRO, DYCLONINE HYDROCHLORIDE
VIVACAINE, BUPIVACAINE HYDROCHLORIDE

SEPTODONT HOLDING

* SEPTODONT HOLDING SAS
ORAVERSE, PHENTOLAMINE MESYLATE

SEPTODONT INC

* SEPTODONT INC
LIDOCAINE, LIDOCAINE
PRILLOCAINE HYDROCHLORIDE AND EPINEPHRINE BITARTRATE, EPINEPHRINE BITARTRATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** s ****

* SEPTODONT INC

PRILOCAINE HYDROCHLORIDE, PRILOCAINE HYDROCHLORIDE

SERVIER

* SERVIER PHARMACEUTICALS LLC

TIBSOVO, IVOSIDENIB

VORANIGO, VORASIDENIB

SETON PHARM

* SETON PHARMACEUTICAL LLC

PEDIAPRED, PREDNISOLONE SODIUM PHOSPHATE

SETON PHARMS

* SETON PHARMACEUTICALS LLC

MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE

PENTAMIDINE ISETHIONATE, PENTAMIDINE ISETHIONATE

SHANDONG

* SHANDONG ANXIN PHARMACEUTICAL CO LTD

PIPERACILLIN AND TAZOBACTAM, PIPERACILLIN SODIUM

* SHANDONG NEW TIME PHARMACEUTICAL CO LTD

ISOSORBIDE MONONITRATE, ISOSORBIDE MONONITRATE

MILRINONE LACTATE, MILRINONE LACTATE

ROCURONIUM BROMIDE, ROCURONIUM BROMIDE

ROSUVASTATIN CALCIUM, ROSUVASTATIN CALCIUM

SEVOFLURANE, SEVOFLURANE

TADALAFIL, TADALAFIL

SHANDONG LUYE

* SHANDONG LUYE PHARMACEUTICAL CO LTD

RYKINDO, RISPERIDONE

SHANDONG XINHUA

* SHANDONG XINHUA PHARMACEUTICAL CO LTD

ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM

IBUPROFEN, IBUPROFEN

IBUPROFEN, IBUPROFEN (OTC)

SEVELAMER CARBONATE, SEVELAMER CARBONATE

SHANGHAI HENGRUI

* SHANGHAI HENGRUI PHARMACEUTICAL CO LTD

DESFLURANE, DESFLURANE

SEVOFLURANE, SEVOFLURANE

SHENZHEN TECHDOW

* SHENZHEN TECHDOW PHARMACEUTICAL CO LTD

ENOXAPARIN SODIUM (PRESERVATIVE FREE), ENOXAPARIN SODIUM

HEPARIN SODIUM PRESERVATIVE FREE, HEPARIN SODIUM

HEPARIN SODIUM, HEPARIN SODIUM

SHIELD TX

* SHIELD TX UK LTD

ACCRUFER, FERRIC MALTOL

SHILPA

* SHILPA MEDICARE LTD

APREMILAST, APREMILAST

BORTEZOMIB, BORTEZOMIB

BUSULFAN, BUSULFAN

CAPECITABINE, CAPECITABINE

DOCETAXEL, DOCETAXEL

ERLOTINIB HYDROCHLORIDE, ERLOTINIB HYDROCHLORIDE

GEMCITABINE HYDROCHLORIDE, GEMCITABINE HYDROCHLORIDE

IMATINIB MESYLATE, IMATINIB MESYLATE

IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE

PEMETREXED, PEMETREXED DISODIUM

SHILPA MEDICARE

* SHILPA MEDICARE LTD

AZACITIDINE, AZACITIDINE

SHIONOGI INC

* SHIONOGI INC

FETROJA, CEFIDEROCOL SULFATE TOSYLATE

SHORLA

* SHORLA PHARMA LTD

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** S ******* SHORLA PHARMA LTD**

JYLAMVO, METHOTREXATE
 NELARABINE, NELARABINE
 TEPYLUTE, THIOTEPA

SHORLA ONCOLOGY*** SHORLA ONCOLOGY**

IMKELDI, IMATINIB MESYLATE

SIGA TECHNOLOGIES*** SIGA TECHNOLOGIES INC**

TPOXX, TECOVIRIMAT

SIGMAPHARM LABS LLC*** SIGMAPHARM LABORATORIES LLC**

ACITRETIN, ACITRETIN
 ADEFOVIR DIPIVOXIL, ADEFOVIR DIPIVOXIL
 AMBRISENTAN, AMBRISENTAN
 AMILORIDE HYDROCHLORIDE, AMILORIDE HYDROCHLORIDE
 ASENAPINE MALEATE, ASENAPINE MALEATE
 DISULFIRAM, DISULFIRAM
 DOFETILIDE, DOFETILIDE
 FLUCYTOSINE, FLUCYTOSINE
 GRISEOFULVIN, GRISEOFULVIN, MICROSIZE
 GRISEOFULVIN, ULTRAMICROSIZE, GRISEOFULVIN, ULTRAMICROSIZE
 LIOTHYRONINE SODIUM, LIOTHYRONINE SODIUM
 PROTRIPTYLINE HYDROCHLORIDE, PROTRIPTYLINE HYDROCHLORIDE
 SODIUM PHENYL BUTYRATE, SODIUM PHENYL BUTYRATE

SINOTHERAPEUTICS INC*** SINOTHERAPEUTICS INC**

BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
 MESALAMINE, MESALAMINE
 POSACONAZOLE, POSACONAZOLE
 PROPAFENONE HYDROCHLORIDE, PROPAFENONE HYDROCHLORIDE

SINTETICA US*** SINTETICA US LLC**

LIDOCAINE HYDROCHLORIDE, LIDOCAINE HYDROCHLORIDE

SK LIFE*** SK LIFE SCIENCE INC**

XCOPRI, CENOBAMATE

SKG PHARMA*** SKG PHARMA INC**

CABERGOLINE, CABERGOLINE
 NICARDIPINE HYDROCHLORIDE, NICARDIPINE HYDROCHLORIDE
 RASAGILINE MESYLATE, RASAGILINE MESYLATE

SLATE RUN PHARMA*** SLATE RUN PHARMACEUTICALS LLC**

ACYCLOVIR SODIUM, ACYCLOVIR SODIUM
 AZITHROMYCIN, AZITHROMYCIN
 BIVALIRUDIN, BIVALIRUDIN
 CILOSTAZOL, CILOSTAZOL
 CINACALCET HYDROCHLORIDE, CINACALCET HYDROCHLORIDE
 DICYCLOMINE HYDROCHLORIDE, DICYCLOMINE HYDROCHLORIDE
 DOBUTAMINE HYDROCHLORIDE, DOBUTAMINE HYDROCHLORIDE
 DOXYCYCLINE HCLATE, DOXYCYCLINE HCLATE
 EPTIFIBATIDE, EPTIFIBATIDE
 ESOMEPRAZOLE SODIUM, ESOMEPRAZOLE SODIUM
 GANCICLOVIR SODIUM, GANCICLOVIR SODIUM
 METHOCARBAMOL, METHOCARBAMOL
 PROMETHAZINE DM, DEXTROMETHORPHAN HYDROBROMIDE
 SODIUM NITROPRUSSIDE, SODIUM NITROPRUSSIDE
 TOBRAMYCIN SULFATE, TOBRAMYCIN SULFATE
 VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE
 VORICONAZOLE, VORICONAZOLE

SOBI*** SOBI INC**

VONJO, PACRITINIB CITRATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** s ******SOFGEN PHARMS**

- * SOFGEN PHARMACEUTICALS LLC
IBUPROFEN, IBUPROFEN (OTC)
OMEGA-3-ACID ETHYL ESTERS, OMEGA-3-ACID ETHYL ESTERS
PROGESTERONE, PROGESTERONE

SOFIE

- * SOFIE CO DBA SOFIE
AMMONIA N 13, AMMONIA N-13
- * SOFIE CO DBA SOFIE (FKA ZEVACOR PHARMA INC)
FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18
SODIUM FLUORIDE F-18, SODIUM FLUORIDE F-18

SOLA PHARMS

- * SOLA PHARMACEUTICALS
CARBOPROST TROMETHAMINE, CARBOPROST TROMETHAMINE
CYANOCOBALAMIN, CYANOCOBALAMIN
DIMETHYL FUMARATE, DIMETHYL FUMARATE
TERIFLUNOMIDE, TERIFLUNOMIDE

SOLARIS PHARMA CORP

- * SOLARIS PHARMA CORP
CLINDAMYCIN PHOSPHATE AND TRETINOIN, CLINDAMYCIN PHOSPHATE
CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
ESTRADIOL, ESTRADIOL
METRONIDAZOLE, METRONIDAZOLE
TAZAROTENE, TAZAROTENE

SOLUBIOMIX

- * SOLUBIOMIX LLC
COXANTO, OXAPROZIN

SOMERSET

- * SOMERSET PHARMACEUTICALS INC
EMSAM, SELEGILINE
- * SOMERSET THERAPEUTICS LLC
ATROPINE SULFATE, ATROPINE SULFATE
BRIMONIDINE TARTRATE AND TIMOLOL MALEATE, BRIMONIDINE TARTRATE
BUPIVACAINE HYDROCHLORIDE, BUPIVACAINE HYDROCHLORIDE
CALCIUM CHLORIDE 10%, CALCIUM CHLORIDE
CALCIUM GLUCONATE, CALCIUM GLUCONATE
CISATRACURIUM BESYLATE, CISATRACURIUM BESYLATE
CUPRIC CHLORIDE, CUPRIC CHLORIDE
DEXAMETHASONE SODIUM PHOSPHATE, DEXAMETHASONE SODIUM PHOSPHATE
DEXMETETOMIDINE HYDROCHLORIDE, DEXMETETOMIDINE HYDROCHLORIDE
DORZOLAMIDE HYDROCHLORIDE AND TIMOLOL MALEATE, DORZOLAMIDE HYDROCHLORIDE
LATANOPROST, LATANOPROST
OFLOXACIN, OFLOXACIN
OLOPATADINE HYDROCHLORIDE, OLOPATADINE HYDROCHLORIDE (OTC)
SULFAMETHOXAZOLE AND TRIMETHOPRIM, SULFAMETHOXAZOLE
TIMOLOL MALEATE, TIMOLOL MALEATE
TIMOLOL, TIMOLOL
VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE
ZINC CHLORIDE, ZINC CHLORIDE

SOMERSET THERAPS LLC

- * SOMERSET THERAPEUTICS LLC
ACARBOSE, ACARBOSE
ACETYLCYSTEINE, ACETYLCYSTEINE
ATROPINE SULFATE, ATROPINE SULFATE
AZELASTINE HYDROCHLORIDE, AZELASTINE HYDROCHLORIDE
BEPOTASTINE BESILATE, BEPOTASTINE BESILATE
BIMATOPROST, BIMATOPROST
BRIMONIDINE TARTRATE, BRIMONIDINE TARTRATE
CABERGOLINE, CABERGOLINE
CISATRACURIUM BESYLATE PRESERVATIVE FREE, CISATRACURIUM BESYLATE
CYANOCOBALAMIN, CYANOCOBALAMIN
DEXAMETHASONE SODIUM PHOSPHATE PRESERVATIVE FREE, DEXAMETHASONE SODIUM PHOSPHATE
DEXAMETHASONE SODIUM PHOSPHATE, DEXAMETHASONE SODIUM PHOSPHATE
DICYCLOMINE HYDROCHLORIDE, DICYCLOMINE HYDROCHLORIDE
DROXIDOPA, DROXIDOPA

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** S ******* SOMERSET THERAPEUTICS LLC**

EPINASTINE HYDROCHLORIDE, EPINASTINE HYDROCHLORIDE
 GLYCOPYRROLATE, GLYCOPYRROLATE
 HALOPERIDOL DECANOATE, HALOPERIDOL DECANOATE
 ISOPROTERENOL HYDROCHLORIDE, ISOPROTERENOL HYDROCHLORIDE
 LACOSAMIDE, LACOSAMIDE
 METHOCARBAMOL, METHOCARBAMOL
 MOXIFLOXACIN HYDROCHLORIDE, MOXIFLOXACIN HYDROCHLORIDE
 NALBUPHINE HYDROCHLORIDE, NALBUPHINE HYDROCHLORIDE
 NALOXONE HYDROCHLORIDE, NALOXONE HYDROCHLORIDE
 OFLOXACIN, OFLOXACIN
 OLOPATADINE HYDROCHLORIDE, OLOPATADINE HYDROCHLORIDE (OTC)
 PILOCARPINE HYDROCHLORIDE, PILOCARPINE HYDROCHLORIDE
 POTASSIUM PHOSPHATES, POTASSIUM PHOSPHATE, DIBASIC
 PROCHLORPERAZINE EDISYLATE, PROCHLORPERAZINE EDISYLATE
 PROPARACAINE HYDROCHLORIDE, PROPARACAINE HYDROCHLORIDE
 ROPIVACAINE HYDROCHLORIDE, ROPIVACAINE HYDROCHLORIDE
 SILDENAFIL CITRATE, SILDENAFIL CITRATE
 SODIUM NITROPRUSSIDE, SODIUM NITROPRUSSIDE
 SUCCINYLCHOLINE CHLORIDE, SUCCINYLCHOLINE CHLORIDE
 TETRACAINE HYDROCHLORIDE, TETRACAINE HYDROCHLORIDE
 TIMOLOL MALEATE, TIMOLOL MALEATE
 TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE
 TOBRAMYCIN, TOBRAMYCIN
 TRAVOPROST, TRAVOPROST
 TRIMETHOPRIM SULFATE AND POLYMYXIN B SULFATE, POLYMYXIN B SULFATE
 TROPICAMIDE, TROPICAMIDE
 VALGANCICLOVIR HYDROCHLORIDE, VALGANCICLOVIR HYDROCHLORIDE
 VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE
 ZINC SULFATE, ZINC SULFATE
 ZOLMITRIPTAN, ZOLMITRIPTAN

SPECGX LLC*** SPECGX LLC**

ACETAMINOPHEN AND CODEINE PHOSPHATE, ACETAMINOPHEN
 AMPHETAMINE SULFATE, AMPHETAMINE SULFATE
 ANAFRANIL, CLOMIPRAMINE HYDROCHLORIDE
 ANEXSIA 5/325, ACETAMINOPHEN
 ANEXSIA 7.5/325, ACETAMINOPHEN
 BUPRENORPHINE HYDROCHLORIDE AND NALOXONE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE
 DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE
 DEXTROAMPHETAMINE SULFATE, DEXTROAMPHETAMINE SULFATE
 DIPHENOXYLATE HYDROCHLORIDE AND ATROPINE SULFATE, ATROPINE SULFATE
 FENTANYL-100, FENTANYL
 FENTANYL-12, FENTANYL
 FENTANYL-25, FENTANYL
 FENTANYL-37, FENTANYL
 FENTANYL-50, FENTANYL
 FENTANYL-62, FENTANYL
 FENTANYL-75, FENTANYL
 HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
 HYDROMORPHONE HYDROCHLORIDE, HYDROMORPHONE HYDROCHLORIDE
 LEVORPHANOL TARTRATE, LEVORPHANOL TARTRATE
 LISDEXAMFETAMINE DIMESYLATE, LISDEXAMFETAMINE DIMESYLATE
 METHADONE HYDROCHLORIDE, METHADONE HYDROCHLORIDE
 METHADOSE, METHADONE HYDROCHLORIDE
 METHYLIN ER, METHYLPHENIDATE HYDROCHLORIDE
 METHYLIN, METHYLPHENIDATE HYDROCHLORIDE
 METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
 MORPHINE SULFATE, MORPHINE SULFATE
 NALTREXONE HYDROCHLORIDE, NALTREXONE HYDROCHLORIDE
 OXYCET, ACETAMINOPHEN
 OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN
 OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
 POSACONAZOLE, POSACONAZOLE
 RESTORIL, TEMAZEPAM

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** S ****

- * SPECGX LLC
ROXICODONE, OXYCODONE HYDROCHLORIDE
- SPECTRA MDCL DEVICES**
- * SPECTRA MEDICAL DEVICES INC
LIDOCAINE HYDROCHLORIDE, LIDOCAINE HYDROCHLORIDE
SODIUM CHLORIDE 0.9%, SODIUM CHLORIDE
- SPIL**
- * SUN PHARMA INDUSTRIES LTD
AMPHOTERICIN B, AMPHOTERICIN B
KAPSPARGO SPRINKLE, METOPROLOL SUCCINATE
NIFEDIPINE, NIFEDIPINE
- SPRIASO LLC**
- * SPRIASO LLC
ICOSAPENT ETHYL, ICOSAPENT ETHYL
- SPRINGWORKS**
- * SPRINGWORKS THERAPEUTICS INC
OGSIVEO, NIROGACESTAT HYDROBROMIDE
- SPROUT PHARMS**
- * SPROUT PHARMACEUTICALS INC
ADDYI, FLIBANSERIN
- SQUARE PHARMS**
- * SQUARE PHARMACEUTICALS LTD
ACYCLOVIR, ACYCLOVIR
ARIPIPIRAZOLE, ARIPIPIRAZOLE
CALCIUM ACETATE, CALCIUM ACETATE
MIRTAZAPINE, MIRTAZAPINE
VALSARTAN, VALSARTAN
- ST RENATUS**
- * ST RENATUS LLC
KOVANAZE, OXYMETAZOLINE HYDROCHLORIDE
- STAND HOMEOPATH**
- * STANDARD HOMEOPATHIC CO
IVY BLOCK, BENTOQUATAM (OTC)
- STASON PHARMS**
- * STASON PHARMACEUTICALS INC
PURINETHOL, MERCAPTOPURINE
- STEMLINE THERAP**
- * STEMLINE THERAPEUTICS INC
ORSERDU, ELACESTRANT HYDROCHLORIDE
- STERIMAX**
- * STERIMAX INC
ETHACRYNATE SODIUM, ETHACRYNATE SODIUM
- STERISCIENCE**
- * STERISCIENCE PTE LTD
FOCINVEZ, FOSAPREPITANT DIMEGLUMINE
METHYLENE BLUE, METHYLENE BLUE
ROCURONIUM BROMIDE, ROCURONIUM BROMIDE
SODIUM BICARBONATE, SODIUM BICARBONATE
SUCCINYLMCHOLINE CHLORIDE, SUCCINYLMCHOLINE CHLORIDE
- STERISCIENCE SPECLTS**
- * STERISCIENCE SPECIALITIES PTE LTD
ACETYLCYSTEINE, ACETYLCYSTEINE
AMPICILLIN AND SULBACTAM, AMPICILLIN SODIUM
AMPICILLIN SODIUM, AMPICILLIN SODIUM
BUPIVACAINE HYDROCHLORIDE PRESERVATIVE FREE, BUPIVACAINE HYDROCHLORIDE
BUPIVACAINE HYDROCHLORIDE, BUPIVACAINE HYDROCHLORIDE
CARBOPROST TROMETHAMINE, CARBOPROST TROMETHAMINE
METHOHEXITAL SODIUM, METHOHEXITAL SODIUM
MIDAZOLAM HYDROCHLORIDE PRESERVATIVE FREE, MIDAZOLAM HYDROCHLORIDE
MIDOZALAM HYDROCHLORIDE, MIDAZOLAM HYDROCHLORIDE
MYCOPHENOLATE MOFETIL HYDROCHLORIDE, MYCOPHENOLATE MOFETIL HYDROCHLORIDE
NAFCILLIN SODIUM, NAFCILLIN SODIUM
OXACILLIN SODIUM, OXACILLIN SODIUM
PIPERACILLIN AND TAZOBACTAM, PIPERACILLIN SODIUM

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** S ****

* STERISCIENCE SPECIALITIES PTE LTD
 SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
 THIAMINE HYDROCHLORIDE, THIAMINE HYDROCHLORIDE

STIEFEL

* STIEFEL LABORATORIES INC
 DUAC, BENZOYL PEROXIDE

STRIDES PHARMA

* STRIDES PHARMA GLOBAL PTE LTD
 ACARBOSE, ACARBOSE
 ACCOLATE, ZAFIRLUKAST
 ACETAMINOPHEN AND CODEINE PHOSPHATE, ACETAMINOPHEN
 ACETAZOLAMIDE, ACETAZOLAMIDE
 ACYCLOVIR, ACYCLOVIR
 ALPRAZOLAM, ALPRAZOLAM
 AMANTADINE HYDROCHLORIDE, AMANTADINE HYDROCHLORIDE
 AMLODIPINE BESYLATE AND VALSARTAN, AMLODIPINE BESYLATE
 AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
 AMLODIPINE BESYLATE, VALSARTAN AND HYDROCHLOROTHIAZIDE, AMLODIPINE BESYLATE
 ATOMOXETINE HYDROCHLORIDE, ATOMOXETINE HYDROCHLORIDE
 AZITHROMYCIN, AZITHROMYCIN
 BUSPIRONE HYDROCHLORIDE, BUSPIRONE HYDROCHLORIDE
 BUTALBITAL, ACETAMINOPHEN AND CAFFEINE, ACETAMINOPHEN
 BUTALBITAL, ACETAMINOPHEN, CAFFEINE AND CODEINE PHOSPHATE, ACETAMINOPHEN
 BUTALBITAL, ASPIRIN AND CAFFEINE, ASPIRIN
 CABERGOLINE, CABERGOLINE
 CINACALCET HYDROCHLORIDE, CINACALCET HYDROCHLORIDE
 COLCHICINE, COLCHICINE
 CYPROHEPTADINE HYDROCHLORIDE, CYPROHEPTADINE HYDROCHLORIDE
 DEXTROAMPHETAMINE SULFATE, DEXTROAMPHETAMINE SULFATE
 DIAZEPAM, DIAZEPAM
 DOFETILIDE, DOFETILIDE
 DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
 DOXEPIN HYDROCHLORIDE, DOXEPIN HYDROCHLORIDE
 DOXYCYCLINE, DOXYCYCLINE
 EFAVIRENZ, EFAVIRENZ
 EMTRICITABINE AND TENOFOVIR DISOPROXIL FUMARATE, EMTRICITABINE
 FLUCYTOSINE, FLUCYTOSINE
 FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 GABAPENTIN, GABAPENTIN
 HYDRA-ZIDE, HYDRALAZINE HYDROCHLORIDE
 HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
 HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
 HYDROCORTISONE, HYDROCORTISONE
 IBUPROFEN, IBUPROFEN
 IBUPROFEN, IBUPROFEN (OTC)
 IMIPRAMINE HYDROCHLORIDE, IMIPRAMINE HYDROCHLORIDE
 KETOCONAZOLE, KETOCONAZOLE
 LAMIVUDINE AND ZIDOVUDINE, LAMIVUDINE
 LAMIVUDINE, LAMIVUDINE
 LEVETIRACETAM, LEVETIRACETAM
 LIDOCAINE, LIDOCAINE
 LOSARTAN POTASSIUM, LOSARTAN POTASSIUM
 MEGESTROL ACETATE, MEGESTROL ACETATE
 MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
 METOCLOPRAMIDE HYDROCHLORIDE, METOCLOPRAMIDE HYDROCHLORIDE
 METRONIDAZOLE, METRONIDAZOLE
 MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE
 MORPHINE SULFATE, MORPHINE SULFATE
 MYCOPHENOLATE MOFETIL, MYCOPHENOLATE MOFETIL
 NATEGLINIDE, NATEGLINIDE
 NEVIRAPINE, NEVIRAPINE
 NYSTATIN AND TRIAMCINOLONE ACETONIDE, NYSTATIN
 OLANZAPINE, OLANZAPINE
 OMEPRAZOLE AND SODIUM BICARBONATE, OMEPRAZOLE
 OSELTAMIVIR PHOSPHATE, OSELTAMIVIR PHOSPHATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** S ****

* STRIDES PHARMA GLOBAL PTE LTD
 OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE
 OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
 PEG-3350, POTASSIUM CHLORIDE, SODIUM BICARBONATE, SODIUM CHLORIDE, POLYETHYLENE GLYCOL
 PIROXICAM, PIROXICAM
 POLYETHYLENE GLYCOL 3350 AND ELECTROLYTES, POLYETHYLENE GLYCOL 3350
 POLYETHYLENE GLYCOL 3350, POLYETHYLENE GLYCOL 3350 (OTC)
 POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
 POTASSIUM CITRATE, POTASSIUM CITRATE
 PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE
 PREDNISONE, PREDNISONE
 PREGABALIN, PREGABALIN
 PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
 PROPAFENONE HYDROCHLORIDE, PROPAFENONE HYDROCHLORIDE
 ROFLUMILAST, ROFLUMILAST
 SERTRALINE HYDROCHLORIDE, SERTRALINE HYDROCHLORIDE
 SEVELAMER CARBONATE, SEVELAMER CARBONATE
 SODIUM SULFATE, POTASSIUM SULFATE AND MAGNESIUM SULFATE, MAGNESIUM SULFATE
 SUCRALFATE, SUCRALFATE
 SUMYCIN, TETRACYCLINE HYDROCHLORIDE
 TACROLIMUS, TACROLIMUS
 TENOFOVIR DISOPROXIL FUMARATE, TENOFOVIR DISOPROXIL FUMARATE
 TESTOSTERONE, TESTOSTERONE
 THEOPHYLLINE, THEOPHYLLINE
 TOPIRAMATE, TOPIRAMATE
 TORSEMIDE, TORSEMIDE
 TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE
 TRANLYCYPROMINE SULFATE, TRANLYCYPROMINE SULFATE
 TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE
 URSODIOL, URSODIOL
 VALGANCICLOVIR HYDROCHLORIDE, VALGANCICLOVIR HYDROCHLORIDE
 VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE
 VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE
 ZILEUTON, ZILEUTON

STRIDES SOFTGELS

* STRIDES SOFTGELS PTE LTD
 AMANTADINE HYDROCHLORIDE, AMANTADINE HYDROCHLORIDE
 BENZONATATE, BENZONATATE
 CALCITRIOL, CALCITRIOL
 CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 CYCLOSPORINE, CYCLOSPORINE
 DICLOFENAC POTASSIUM, DICLOFENAC POTASSIUM
 DUTASTERIDE, DUTASTERIDE
 ERGOCALCIFEROL, ERGOCALCIFEROL
 ETHOSUXIMIDE, ETHOSUXIMIDE
 IBUPROFEN AND DIPHENHYDRAMINE HYDROCHLORIDE, DIPHENHYDRAMINE HYDROCHLORIDE (OTC)
 IBUPROFEN, IBUPROFEN (OTC)
 ICOSAPENT ETHYL, ICOSAPENT ETHYL
 LOPERAMIDE HYDROCHLORIDE, LOPERAMIDE HYDROCHLORIDE (OTC)
 METHOXSALEN, METHOXSALEN
 OMEGA-3-ACID ETHYL ESTERS, OMEGA-3-ACID ETHYL ESTERS

STRONGBRIDGE

* STRONGBRIDGE DUBLIN LTD
 RECORLEV, LEVOKETOCONAZOLE

SUCAMPO PHARMA LLC

* SUCAMPO PHARMA AMERICAS LLC
 AMITIZA, LUBIPROSTONE

SUMITOMO PHARMA

* SUMITOMO PHARMA SWITZERLAND GMBH
 MYFEMBREE, ESTRADIOL
 ORGOVYX, RELUGOLIX

SUMITOMO PHARMA AM

* SUMITOMO PHARMA AMERICA INC
 APTIOM, ESLICARBAZEPINE ACETATE

SUMMIT BIOSCI

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** S ****

* SUMMIT BIOSCIENCES LLC

REZENOPY, NALOXONE HYDROCHLORIDE

SUN PHARM

* SUN PHARMACEUTICAL INDUSTRIES LTD

ABSORICA LD, ISOTRETINOIN

ALBUTEROL SULFATE AND IPRATROPIUM BROMIDE, ALBUTEROL SULFATE

ALBUTEROL SULFATE, ALBUTEROL SULFATE

ALENDRONATE SODIUM, ALENDRONATE SODIUM

ALPRAZOLAM, ALPRAZOLAM

AMBRISENTAN, AMBRISENTAN

ARFORMOTEROL TARTRATE, ARFORMOTEROL TARTRATE

BICALUTAMIDE, BICALUTAMIDE

BOSENTAN, BOSENTAN

BROMSITE, BROMFENAC SODIUM

BUDESONIDE, BUDESONIDE

BUPRENORPHINE HYDROCHLORIDE AND NALOXONE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE

BUPRENORPHINE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE

BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE

BYNFEZIA PEN, OCTREOTIDE ACETATE

CAFFEINE CITRATE, CAFFEINE CITRATE

CARBIDOPA AND LEVODOPA, CARBIDOPA

CARBIDOPA, LEVODOPA AND ENTACAPONE, CARBIDOPA

CARBOPLATIN, CARBOPLATIN

CEQUA, CYCLOSPORINE

CHLORPROMAZINE HYDROCHLORIDE, CHLORPROMAZINE HYDROCHLORIDE

CINACALCET HYDROCHLORIDE, CINACALCET HYDROCHLORIDE

CIPROFLOXACIN AND DEXAMETHASONE, CIPROFLOXACIN

DALFAMPRIDINE, DALFAMPRIDINE

DECITABINE, DECITABINE

DEFERASIROX, DEFERASIROX

DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE

DOFETILIDE, DOFETILIDE

DOXORUBICIN HYDROCHLORIDE (LIPOSOMAL), DOXORUBICIN HYDROCHLORIDE

DRIZALMA SPRINKLE, DULOXETINE HYDROCHLORIDE

DROXIDOPA, DROXIDOPA

ENTACAPONE, ENTACAPONE

EPOPROSTENOL SODIUM, EPOPROSTENOL SODIUM

ERLOTINIB HYDROCHLORIDE, ERLOTINIB HYDROCHLORIDE

ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM (OTC)

ESZOPICLONE, ESZOPICLONE

FEBUXOSTAT, FEBUXOSTAT

FENOFIBRATE, FENOFIBRATE

FEXOFENADINE HYDROCHLORIDE AND PSEUDOEPHEDRINE HYDROCHLORIDE, FEXOFENADINE

FINASTERIDE, FINASTERIDE

FOSAMPRENAVIR CALCIUM, FOSAMPRENAVIR CALCIUM

FOSPHENYTOIN SODIUM, FOSPHENYTOIN SODIUM

FYREMADEL, GANIRELIX ACETATE

GALANTAMINE HYDROBROMIDE, GALANTAMINE HYDROBROMIDE

GEMCITABINE HYDROCHLORIDE, GEMCITABINE HYDROCHLORIDE

GUAIFENESIN AND DEXTROMETHORPHAN HYDROBROMIDE, DEXTROMETHORPHAN HYDROBROMIDE (OTC)

GUANFACINE HYDROCHLORIDE, GUANFACINE HYDROCHLORIDE

IMATINIB MESYLATE, IMATINIB MESYLATE

IPRATROPIUM BROMIDE, IPRATROPIUM BROMIDE

KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE

LACOSAMIDE, LACOSAMIDE

LENALIDOMIDE, LENALIDOMIDE

LEUPROLIDE ACETATE, LEUPROLIDE ACETATE

LEVOCETIRIZINE DIHYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE

LIOTHYRONINE SODIUM, LIOTHYRONINE SODIUM

LORATADINE, LORATADINE (OTC)

LOTEPREDNOL ETABONATE, LOTEPREDNOL ETABONATE

LURASIDONE HYDROCHLORIDE, LURASIDONE HYDROCHLORIDE

MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE

MESALAMINE, MESALAMINE

METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** S ****

* SUN PHARMACEUTICAL INDUSTRIES LTD
 METHOTREXATE SODIUM, METHOTREXATE SODIUM
 MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE
 NALTREXONE HYDROCHLORIDE, NALTREXONE HYDROCHLORIDE
 NIACIN, NIACIN
 ODOMZO, SONIDEGIB PHOSPHATE
 OMEGA-3-ACID ETHYL ESTERS, OMEGA-3-ACID ETHYL ESTERS
 OMEPRAZOLE, OMEPRAZOLE (OTC)
 OPCICON ONE-STEP, LEVONORGESTREL (OTC)
 PALIPERIDONE, PALIPERIDONE
 PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
 PAZOPANIB HYDROCHLORIDE, PAZOPANIB HYDROCHLORIDE
 QUETIAPINE FUMARATE, QUETIAPINE FUMARATE
 RANOLAZINE, RANOLAZINE
 RISEDRONATE SODIUM, RISEDRONATE SODIUM
 SAXAGLIPTIN AND METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 SAXAGLIPTIN, SAXAGLIPTIN HYDROCHLORIDE
 SUMATRIPTAN AND NAPROXEN SODIUM, NAPROXEN SODIUM
 SUNITINIB MALATE, SUNITINIB MALATE
 TADALAFIL, TADALAFIL
 TEMOZOLOMIDE, TEMOZOLOMIDE
 TOBRAMYCIN, TOBRAMYCIN
 TOPIRAMATE, TOPIRAMATE
 VECURONIUM BROMIDE, VECURONIUM BROMIDE
 WINLEVI, CLASCOTERONE
 YONSA, ABIRATERONE ACETATE
 ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE

SUN PHARM INDS

* SUN PHARMACEUTICAL INDUSTRIES LTD
 CARBIDOPA AND LEVODOPA, CARBIDOPA
 DIVALPROEX SODIUM, DIVALPROEX SODIUM
 DOXORUBICIN HYDROCHLORIDE, DOXORUBICIN HYDROCHLORIDE
 FEXOFENADINE HYDROCHLORIDE ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
 FEXOFENADINE HYDROCHLORIDE HIVES, FEXOFENADINE HYDROCHLORIDE (OTC)
 ONDANSETRON, ONDANSETRON
 OXCARBAZEPINE, OXCARBAZEPINE
 SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
 TIAGABINE HYDROCHLORIDE, TIAGABINE HYDROCHLORIDE

SUN PHARM INDS (IN)

* SUN PHARMACEUTICAL INDUSTRIES LTD
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 ZONISAMIDE, ZONISAMIDE

SUN PHARM INDS INC

* SUN PHARMACEUTICAL INDUSTRIES INC
 ABSORICA, ISOTRETINOIN
 AMITRIPTYLINE HYDROCHLORIDE, AMITRIPTYLINE HYDROCHLORIDE
 AN-SULFUR COLLOID, TECHNETIUM TC-99M SULFUR COLLOID KIT
 CIS-MDP, TECHNETIUM TC-99M MEDRONATE KIT
 CIS-PYRO, TECHNETIUM TC-99M PYROPHOSPHATE KIT
 CLONAZEPAM, CLONAZEPAM
 CLOZAPINE, CLOZAPINE
 DEXMETHYLPHENIDATE HYDROCHLORIDE, DEXMETHYLPHENIDATE HYDROCHLORIDE
 DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE
 DIGOXIN, DIGOXIN
 GUAIFENESIN AND PSEUDOEPHEDRINE HYDROCHLORIDE, GUAIFENESIN (OTC)
 HALOG, HALCINONIDE
 INDIUM IN-111 PENTETREOTIDE KIT, INDIUM IN-111 PENTETREOTIDE KIT
 LEQSELVI, DEURUXOLITINIB PHOSPHATE
 LEVORPHANOL TARTRATE, LEVORPHANOL TARTRATE
 LEVULAN, AMINOLEVULINIC ACID HYDROCHLORIDE
 LISDEXAMFETAMINE DIMESYLATE, LISDEXAMFETAMINE DIMESYLATE
 LITHIUM CARBONATE, LITHIUM CARBONATE
 METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
 METOPROLOL TARTRATE, METOPROLOL TARTRATE
 MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** S ****

* SUN PHARMACEUTICAL INDUSTRIES INC
 MIRTAZAPINE, MIRTAZAPINE
 REPAGLINIDE, REPAGLINIDE
 SEZABY, PHENOBARBITAL SODIUM
 TECHNETIUM TC 99M SESTAMIBI, TECHNETIUM TC-99M SESTAMIBI KIT
 TECHNETIUM TC-99M MEBROFENIN, TECHNETIUM TC-99M MEBROFENIN KIT
 TECHNETIUM TC99M MERTIATIDE KIT, TECHNETIUM TC-99M MERTIATIDE KIT
 TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE
 TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE
 VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE

SUN PHARM INDS LTD

* SUN PHARMACEUTICAL INDUSTRIES LTD
 ACETAMINOPHEN AND CODEINE PHOSPHATE, ACETAMINOPHEN
 ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM
 BENAZEPRIL HYDROCHLORIDE, BENAZEPRIL HYDROCHLORIDE
 CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 CETIRIZINE HYDROCHLORIDE AND PSEUDOEPHEDRINE HYDROCHLORIDE, CETIRIZINE HYDROCHLORIDE
 CETIRIZINE HYDROCHLORIDE HIVES, CETIRIZINE HYDROCHLORIDE (OTC)
 CLINDAMYCIN HYDROCHLORIDE, CLINDAMYCIN HYDROCHLORIDE
 CYCLOBENZAPRINE HYDROCHLORIDE, CYCLOBENZAPRINE HYDROCHLORIDE
 DOXYCYCLINE, DOXYCYCLINE
 FELODIPINE, FELODIPINE
 FENOFIBRATE, FENOFIBRATE
 FLECAINIDE ACETATE, FLECAINIDE ACETATE
 GLYCOPYRROLATE, GLYCOPYRROLATE
 LISINAPRIL AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 LISINAPRIL, LISINAPRIL
 LORATADINE AND PSEUDOEPHEDRINE SULFATE, LORATADINE (OTC)
 LORATADINE REDIDOSE, LORATADINE (OTC)
 LORATADINE, LORATADINE (OTC)
 LORAZEPAM, LORAZEPAM
 MORPHINE SULFATE, MORPHINE SULFATE
 NALOXONE HYDROCHLORIDE AND PENTAZOCINE HYDROCHLORIDE, NALOXONE HYDROCHLORIDE
 OXCARBAZEPINE, OXCARBAZEPINE
 PSEUDOEPHEDRINE HYDROCHLORIDE, PSEUDOEPHEDRINE HYDROCHLORIDE (OTC)
 RISPERIDONE, RISPERIDONE
 VALACYCLOVIR HYDROCHLORIDE, VALACYCLOVIR HYDROCHLORIDE

SUN PHARM INDUSTRIES

* SUN PHARMACEUTICAL INDUSTRIES INC
 ALBUTEROL SULFATE, ALBUTEROL SULFATE
 ALLOPURINOL, ALLOPURINOL
 BACTRIM DS, SULFAMETHOXAZOLE
 BACTRIM, SULFAMETHOXAZOLE
 CARVEDILOL PHOSPHATE, CARVEDILOL PHOSPHATE
 CHLOROTHALIDONE, CHLOROTHALIDONE
 DEXMETHYLPHENIDATE HYDROCHLORIDE, DEXMETHYLPHENIDATE HYDROCHLORIDE
 DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE
 DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
 IMIPRAMINE HYDROCHLORIDE, IMIPRAMINE HYDROCHLORIDE
 MINOXIDIL, MINOXIDIL
 NITROFURANTOIN, NITROFURANTOIN, MACROCRYSTALLINE
 PHENTERMINE HYDROCHLORIDE, PHENTERMINE HYDROCHLORIDE
 PINDOLOL, PINDOLOL
 PREDNISONE, PREDNISONE
 QUINIDINE GLUCONATE, QUINIDINE GLUCONATE
 SPIRONOLACTONE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 SPIRONOLACTONE, SPIRONOLACTONE
 SULFAMETHOXAZOLE AND TRIMETHOPRIM, SULFAMETHOXAZOLE
 SULINDAC, SULINDAC
 TEMAZEPAM, TEMAZEPAM
 TRAZODONE HYDROCHLORIDE, TRAZODONE HYDROCHLORIDE

SUNNY

* SUNNY PHARMTECH INC
 AMINOCAPROIC ACID, AMINOCAPROIC ACID
 CARBOPROST TROMETHAMINE, CARBOPROST TROMETHAMINE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** S ****

* SUNNY PHARMTECH INC
CYCLOPHOSPHAMIDE, CYCLOPHOSPHAMIDE
NITROFURANTOIN (MONOHYDRATE/MACROCRYSTALS), NITROFURANTOIN

SUNOVION PHARMS INC

* SUNOVION PHARMACEUTICALS INC
LATUDA, LURASIDONE HYDROCHLORIDE

SUNSHINE

* SUNSHINE LAKE PHARMA CO LTD
ARIPIPIRAZOLE, ARIPIPIRAZOLE
AZITHROMYCIN, AZITHROMYCIN
CLARITHROMYCIN, CLARITHROMYCIN
DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE
ENTACAPONE, ENTACAPONE
FEBUXOSTAT, FEBUXOSTAT
LINAGLIPTIN AND METFORMIN HYDROCHLORIDE, LINAGLIPTIN
LINAGLIPTIN, LINAGLIPTIN
METOPROLOL SUCCINATE, METOPROLOL SUCCINATE
OLMESARTAN MEDOXOMIL, OLMESARTAN MEDOXOMIL
OSELTAMIVIR PHOSPHATE, OSELTAMIVIR PHOSPHATE
RANOLAZINE, RANOLAZINE
SILDENAFIL CITRATE, SILDENAFIL CITRATE
TADALAFIL, TADALAFIL

SUPERNUS PHARMS

* SUPERNUS PHARMACEUTICALS INC
GOCOVRI, AMANTADINE HYDROCHLORIDE
OXTELLAR XR, OXCARBAZEPINE
QELBREE, VILOXAZINE HYDROCHLORIDE
TROKENDI XR, TOPIRAMATE

SUVEN PHARMS

* SUVEN PHARMACEUTICALS LTD
CALCIUM ACETATE, CALCIUM ACETATE
CHLOROQUINE PHOSPHATE, CHLOROQUINE PHOSPHATE
GLYCOPYRROLATE, GLYCOPYRROLATE
MALATHION, MALATHION

SWEDISH ORPHAN

* SWEDISH ORPHAN BIOVITRUM AB PUBL
ORFADIN, NITISINONE

SYNDAX

* SYNDAX PHARMACEUTICALS INC
REVUFORJ, REVUMENIB CITRATE

SYNTHON BV

* SYNTHON BV
TAMSULOSIN HYDROCHLORIDE, TAMSULOSIN HYDROCHLORIDE

SYNTHON PHARMS

* SYNTHON PHARMACEUTICALS INC
LEVOCETIRIZINE DIHYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE

SYNTHON PHARMS INC

* SYNTHON PHARMACEUTICALS INC
GLATIRAMER ACETATE, GLATIRAMER ACETATE

**** T ******ACME LABS**

* THE ACME LABORATORIES LTD
CLOPIDOGREL BISULFATE, CLOPIDOGREL BISULFATE
ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE

GEN HOSP

* THE GENERAL HOSPITAL CORP
AMMONIA N 13, AMMONIA N-13

METHODIST

* THE METHODIST HOSP RESEARCH INSTITUTE
AMMONIA N 13, AMMONIA N-13

METHODIST HOSP RES

* THE METHODIST HOSP RESEARCH INSTITUTE
FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** T ******RITEDOSE CORP**

* THE RITEDOSE CORP
 ALBUTEROL SULFATE AND IPRATROPIUM BROMIDE, ALBUTEROL SULFATE
 ALBUTEROL SULFATE, ALBUTEROL SULFATE
 ARFORMOTEROL TARTRATE, ARFORMOTEROL TARTRATE
 FORMOTEROL FUMARATE, FORMOTEROL FUMARATE
 IPRATROPIUM BROMIDE, IPRATROPIUM BROMIDE
 LEVALBUTEROL HYDROCHLORIDE, LEVALBUTEROL HYDROCHLORIDE

TAGI

* TAGI PHARMA INC
 CHOLESTYRAMINE LIGHT, CHOLESTYRAMINE
 CHOLESTYRAMINE, CHOLESTYRAMINE
 DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE
 METHAZOLAMIDE, METHAZOLAMIDE

TAIHO ONCOLOGY

* TAIHO ONCOLOGY INC
 LONSURF, TIPIRACIL HYDROCHLORIDE
 LYTGOBI, FUTIBATINIB

TAKEDA PHARMS USA

* TAKEDA PHARMACEUTICALS USA INC
 ACTOPLUS MET, METFORMIN HYDROCHLORIDE
 ACTOS, PIOGLITAZONE HYDROCHLORIDE
 ADDERALL XR 10, AMPHETAMINE ASPARTATE
 ADDERALL XR 15, AMPHETAMINE ASPARTATE
 ADDERALL XR 20, AMPHETAMINE ASPARTATE
 ADDERALL XR 25, AMPHETAMINE ASPARTATE
 ADDERALL XR 30, AMPHETAMINE ASPARTATE
 ADDERALL XR 5, AMPHETAMINE ASPARTATE
 AGRYLIN, ANAGRELIDE HYDROCHLORIDE
 ALUNBRIG, BRIGATINIB
 CARBATROL, CARBAMAZEPINE
 DEXILANT, DEXLANSOPRAZOLE
 DUETACT, GLIMEPIRIDE
 EOHILIA, BUDESONIDE
 FIRAZYR, ICATIBANT ACETATE
 FOSRENOL, LANTHANUM CARBONATE
 FRUZAQLA, FRUQUINTINIB
 GATTEX KIT, TEDUGLUTIDE
 ICLUSIG, PONATINIB HYDROCHLORIDE
 INTUNIV, GUANFACINE HYDROCHLORIDE
 KAZANO, ALOGLIPTIN BENZOATE
 LIALDA, MESALAMINE
 LIVTENCITY, MARIBAVIR
 MOTEGRITY, PRUCALOPRIDE SUCCINATE
 MYDAYIS, AMPHETAMINE ASPARTATE
 NESINA, ALOGLIPTIN BENZOATE
 NINLARO, IXAZOMIB CITRATE
 OSENI, ALOGLIPTIN BENZOATE
 PENTASA, MESALAMINE
 PREVACID, LANSOPRAZOLE
 ROZEREM, RAMELTEON
 TRINTELLIX, VORTIOXETINE HYDROBROMIDE
 ULORIC, FEBUXOSTAT
 VELCADE, BORTEZOMIB
 VYVANSE, LISDEXAMFETAMINE DIMESYLATE

TAMARANG

* TAMARANG SA
 ROCURONIUM BROMIDE, ROCURONIUM BROMIDE

TARO

* TARO PHARMACEUTICAL INDUSTRIES LTD
 ACETAMINOPHEN, ACETAMINOPHEN (OTC)
 ACETAZOLAMIDE, ACETAZOLAMIDE
 AMINOCAPROIC ACID, AMINOCAPROIC ACID
 AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE
 BETAMETHASONE VALERATE, BETAMETHASONE VALERATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** T ****

- * TARO PHARMACEUTICAL INDUSTRIES LTD
 BROMPHENIRAMINE MALEATE, PSEUDOEPHEDRINE HYDROCHLORIDE AND DEXTROMETHORPHAN
 BUMETANIDE, BUMETANIDE
 BUTALBITAL, ACETAMINOPHEN AND CAFFEINE, ACETAMINOPHEN
 CARBAMAZEPINE, CARBAMAZEPINE
 CARVEDILOL, CARVEDILOL
 CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 CETIRIZINE HYDROCHLORIDE HIVES RELIEF, CETIRIZINE HYDROCHLORIDE (OTC)
 CETIRIZINE HYDROCHLORIDE, CETIRIZINE HYDROCHLORIDE
 CHILDREN'S CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 CHILDREN'S CETIRIZINE HYDROCHLORIDE HIVES RELIEF, CETIRIZINE HYDROCHLORIDE (OTC)
 CHILDREN'S FEXOFENADINE HYDROCHLORIDE ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
 CHILDREN'S FEXOFENADINE HYDROCHLORIDE HIVES, FEXOFENADINE HYDROCHLORIDE (OTC)
 CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
 CLOBAZAM, CLOBAZAM
 CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
 CLOCORTOLONE PIVALATE, CLOCORTOLONE PIVALATE
 CLOMIPRAMINE HYDROCHLORIDE, CLOMIPRAMINE HYDROCHLORIDE
 CLORAZEPATE DIPOTASSIUM, CLORAZEPATE DIPOTASSIUM
 CLOTRIMAZOLE AND BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
 DEFERIPRONE, DEFERIPRONE
 DESONIDE, DESONIDE
 DICLOFENAC POTASSIUM, DICLOFENAC POTASSIUM
 DOXEPIN HYDROCHLORIDE, DOXEPIN HYDROCHLORIDE
 ENALAPRIL MALEATE, ENALAPRIL MALEATE
 ETODOLAC, ETODOLAC
 EXTENDED PHENYTOIN SODIUM, PHENYTOIN SODIUM
 FELBAMATE, FELBAMATE
 FLUCONAZOLE, FLUCONAZOLE
 FLUOCINONIDE EMULSIFIED BASE, FLUOCINONIDE
 FLUOROURACIL, FLUOROURACIL
 FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 FLUPHENAZINE HYDROCHLORIDE, FLUPHENAZINE HYDROCHLORIDE
 GABAPENTIN, GABAPENTIN
 GRANISETRON HYDROCHLORIDE, GRANISETRON HYDROCHLORIDE
 HYDROCORTISONE AND ACETIC ACID, ACETIC ACID, GLACIAL
 HYDROCORTISONE BUTYRATE, HYDROCORTISONE BUTYRATE
 HYDROCORTISONE, HYDROCORTISONE
 IMIQUIMOD, IMIQUIMOD
 INFANTS' FEVERALL, ACETAMINOPHEN (OTC)
 IVERMECTIN, IVERMECTIN (OTC)
 KETOCONAZOLE, KETOCONAZOLE
 LACTULOSE, LACTULOSE
 LAMOTRIGINE, LAMOTRIGINE
 LEVETIRACETAM, LEVETIRACETAM
 LEVOCETIRIZINE DIHYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE
 LORATADINE, LORATADINE (OTC)
 MELOXICAM, MELOXICAM
 METRONIDAZOLE, METRONIDAZOLE
 MINOXIDIL (FOR MEN), MINOXIDIL (OTC)
 MINOXIDIL (FOR WOMEN), MINOXIDIL (OTC)
 NORTRIPTYLINE HYDROCHLORIDE, NORTRIPTYLINE HYDROCHLORIDE
 NYSTATIN, NYSTATIN
 OXCARBAZEPINE, OXCARBAZEPINE
 PEG-3350, SODIUM SULFATE, SODIUM CHLORIDE, POTASSIUM CHLORIDE, SODIUM ASCORBATE AND
 PHENYTOIN, PHENYTOIN
 SILDENAFIL CITRATE, SILDENAFIL CITRATE
 SODIUM SULFATE, POTASSIUM SULFATE AND MAGNESIUM SULFATE, MAGNESIUM SULFATE
 TERIL, CARBAMAZEPINE
 TOPICORT, DESOXIMETASONE
 TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE
 WARFARIN SODIUM, WARFARIN SODIUM
- * TARO PHARMACEUTICALS INC
 ACETIC ACID, ACETIC ACID, GLACIAL
 ACYCLOVIR, ACYCLOVIR

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** T ****

- * TARO PHARMACEUTICALS INC
 ADAPALENE AND BENZOYL PEROXIDE, ADAPALENE
 ADAPALENE, ADAPALENE
 ADAPALENE, ADAPALENE (OTC)
 ALCLOMETASONE DIPROPIONATE, ALCLOMETASONE DIPROPIONATE
 AMMONIUM LACTATE, AMMONIUM LACTATE
 AZELAIC ACID, AZELAIC ACID
 BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
 BUTENAFINE HYDROCHLORIDE, BUTENAFINE HYDROCHLORIDE (OTC)
 CALCIPOTRIENE AND BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
 CICLOPIROX, CICLOPIROX
 CLINDAMYCIN PHOSPHATE AND BENZOYL PEROXIDE, BENZOYL PEROXIDE
 CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
 CLOBETASOL PROPIONATE (EMOLLIENT), CLOBETASOL PROPIONATE
 CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
 CLOTRIMAZOLE AND BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
 CLOTRIMAZOLE, CLOTRIMAZOLE
 CLOTRIMAZOLE, CLOTRIMAZOLE (OTC)
 DAPSONE, DAPSONE
 DERMABET, BETAMETHASONE VALERATE
 DESONIDE, DESONIDE
 DICLOFENAC SODIUM, DICLOFENAC SODIUM
 DIFLORASONE DIACETATE, DIFLORASONE DIACETATE
 DOCOSANOL, DOCOSANOL (OTC)
 ECONAZOLE NITRATE, ECONAZOLE NITRATE
 ESCITALOPRAM OXALATE, ESCITALOPRAM OXALATE
 FLUOCINOLONE ACETONIDE, FLUOCINOLONE ACETONIDE
 FLUOCINONIDE, FLUOCINONIDE
 GENTAMICIN SULFATE, GENTAMICIN SULFATE
 HALOBETASOL PROPIONATE, HALOBETASOL PROPIONATE
 HYDROCORTISONE VALERATE, HYDROCORTISONE VALERATE
 HYDROCORTISONE, HYDROCORTISONE
 IBUPROFEN, IBUPROFEN
 IBUPROFEN, IBUPROFEN (OTC)
 KETOZOLE, KETOCONAZOLE
 LIDOCAINE HYDROCHLORIDE, LIDOCAINE HYDROCHLORIDE
 LIDOCAINE, LIDOCAINE
 LORATADINE, LORATADINE (OTC)
 MICONAZOLE 3, MICONAZOLE NITRATE (OTC)
 MICONAZOLE NITRATE, MICONAZOLE NITRATE (OTC)
 MOMETASONE FUROATE, MOMETASONE FUROATE
 MUPIROCIN, MUPIROCIN
 MUPIROCIN, MUPIROCIN CALCIUM
 NAFTIFINE HYDROCHLORIDE, NAFTIFINE HYDROCHLORIDE
 NYSTATIN AND TRIAMCINOLONE ACETONIDE, NYSTATIN
 NYSTATIN, NYSTATIN
 OXICONAZOLE NITRATE, OXICONAZOLE NITRATE
 PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
 RISPERIDONE, RISPERIDONE
 SULFACETAMIDE SODIUM, SULFACETAMIDE SODIUM
 TAVABOROLE, TAVABOROLE
 TAZAROTENE, TAZAROTENE
 TERBINAFINE HYDROCHLORIDE, TERBINAFINE HYDROCHLORIDE (OTC)
 TERCONAZOLE, TERCONAZOLE
 TOPICORT, DESOXIMETASONE
 TRETINOLIN, TRETINOLIN
 TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE
 TRIVAGIZOLE 3, CLOTRIMAZOLE (OTC)
- * TARO PHARMACEUTICALS USA INC
 SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, SODIUM CHLORIDE
 STERILE WATER FOR INJECTION IN PLASTIC CONTAINER, STERILE WATER FOR INJECTION

TARO PHARM INDS

- * TARO PHARMACEUTICAL INDUSTRIES LTD
 AMCINONIDE, AMCINONIDE
 CARBAMAZEPINE, CARBAMAZEPINE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** T ****

* TARO PHARMACEUTICAL INDUSTRIES LTD
 CICLOPIROX, CICLOPIROX
 CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
 ENALAPRIL MALEATE AND HYDROCHLOROTHIAZIDE, ENALAPRIL MALEATE
 ETODOLAC, ETODOLAC
 HYDROCORTISONE BUTYRATE, HYDROCORTISONE BUTYRATE
 LAMOTRIGINE, LAMOTRIGINE

TARSUS

* TARSUS PHARMACEUTICALS INC
 XDEMVY, LOTILANER

TEIKOKU PHARMA USA

* TEIKOKU PHARMA USA INC
 LIDODERM, LIDOCAINE

TELIGENT

* TELIGENT PHARMA INC
 FLURANDRENOLIDE, FLURANDRENOLIDE

TELIX

* TELIX PHARMACEUTICALS US INC
 ILLUCCIX, GALLIUM GA-68 GOZETOTIDE

TENSHI

* TENSHI KAIZEN PVT LTD
 LORATADINE, LORATADINE (OTC)

TERSERA

* TERSERA THERAPEUTICS LLC
 PRIALT, ZICONOTIDE ACETATE
 VARUBI, ROLAPITANT HYDROCHLORIDE
 XERMELO, TELOTTRISTAT ETIPRATE
 ZOLADEX, GOSERELIN ACETATE

TETRAPHASE PHARMS

* TETRAPHASE PHARMACEUTICALS INC
 XERAVA, ERAVACYCLINE DIHYDROCHLORIDE

TEVA

* TEVA NEUROSCIENCE INC
 AUSTEDO XR, DEUTETRABENAZINE
 AZILECT, RASAGILINE MESYLATE
 UZEDY, RISPERIDONE

* TEVA PHARMACEUTICALS USA INC
 ACYCLOVIR, ACYCLOVIR
 ADIPEX-P, PHENTERMINE HYDROCHLORIDE
 ALBUTEROL SULFATE, ALBUTEROL SULFATE
 AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
 AMOXICILLIN AND CLAVULANATE POTASSIUM, AMOXICILLIN
 AMOXICILLIN PEDIATRIC, AMOXICILLIN
 AMOXICILLIN, AMOXICILLIN
 ATENOLOL, ATENOLOL
 AZITHROMYCIN, AZITHROMYCIN
 BENAZEPRIL HYDROCHLORIDE, BENAZEPRIL HYDROCHLORIDE
 BUSPIRONE HYDROCHLORIDE, BUSPIRONE HYDROCHLORIDE
 CALCITRIOL, CALCITRIOL
 CARVEDILOL, CARVEDILOL
 CEFACLOR, CEFACLOR
 CEFPROZIL, CEFPROZIL
 CELECOXIB, CELECOXIB
 CEPHALEXIN, CEPHALEXIN
 CHILDREN'S FEXOFENADINE HYDROCHLORIDE ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
 CHILDREN'S FEXOFENADINE HYDROCHLORIDE HIVES, FEXOFENADINE HYDROCHLORIDE (OTC)
 CILOSTAZOL, CILOSTAZOL
 CIMETIDINE, CIMETIDINE
 CLONAZEPAM, CLONAZEPAM
 CLOPIDOGREL BISULFATE, CLOPIDOGREL BISULFATE
 CLOTRIMAZOLE, CLOTRIMAZOLE
 DICLOFENAC POTASSIUM, DICLOFENAC POTASSIUM
 DICLOXACILLIN SODIUM, DICLOXACILLIN SODIUM
 DIFLUNISAL, DIFLUNISAL
 DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** T ****

* TEVA PHARMACEUTICALS USA INC
DISOPYRAMIDE PHOSPHATE, DISOPYRAMIDE PHOSPHATE
DOXAZOSIN MESYLATE, DOXAZOSIN MESYLATE
EPITOL, CARBAMAZEPINE
ESZOPICLONE, ESZOPICLONE
ETODOLAC, ETODOLAC
FEXOFENADINE HYDROCHLORIDE ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
FEXOFENADINE HYDROCHLORIDE HIVES, FEXOFENADINE HYDROCHLORIDE (OTC)
FEXOFENADINE HYDROCHLORIDE, FEXOFENADINE HYDROCHLORIDE
FINASTERIDE, FINASTERIDE
FLUOCINONIDE EMULSIFIED BASE, FLUOCINONIDE
FLUOCINONIDE, FLUOCINONIDE
FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
FOSINOPRIL SODIUM, FOSINOPRIL SODIUM
GALZIN, ZINC ACETATE
GLYBURIDE (MICRONIZED), GLYBURIDE
GLYBURIDE, GLYBURIDE
IRBESARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
KETOCONAZOLE, KETOCONAZOLE
KETOPROFEN, KETOPROFEN
KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
LEVOFLOXACIN, LEVOFLOXACIN
LOPERAMIDE HYDROCHLORIDE, LOPERAMIDE HYDROCHLORIDE
LOVASTATIN, LOVASTATIN
METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
METOCLOPRAMIDE HYDROCHLORIDE, METOCLOPRAMIDE HYDROCHLORIDE
MEXILETINE HYDROCHLORIDE, MEXILETINE HYDROCHLORIDE
MIRTAZAPINE, MIRTAZAPINE
MOEXIPRIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
MOEXIPRIL HYDROCHLORIDE, MOEXIPRIL HYDROCHLORIDE
MUPIROCIN, MUPIROCIN
NAPROXEN, NAPROXEN
NEFAZODONE HYDROCHLORIDE, NEFAZODONE HYDROCHLORIDE
NEOMYCIN SULFATE, NEOMYCIN SULFATE
NORTRIPTYLINE HYDROCHLORIDE, NORTRIPTYLINE HYDROCHLORIDE
NYSTATIN, NYSTATIN
OFLOXACIN, OFLOXACIN
OXYMORPHONE HYDROCHLORIDE, OXYMORPHONE HYDROCHLORIDE
PENICILLIN-VK, PENICILLIN V POTASSIUM
PIROXICAM, PIROXICAM
PRELONE, PREDNISOLONE
SILDENAFIL CITRATE, SILDENAFIL CITRATE
SOTALOL HYDROCHLORIDE, SOTALOL HYDROCHLORIDE
SUCRALFATE, SUCRALFATE
TOPIRAMATE, TOPIRAMATE
TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE
VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE
ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE

TEVA BRANDED PHARM

* TEVA BRANDED PHARMACEUTICAL PRODUCTS R AND D INC
ACTIGALL, URSODIOL
AUSTEDO, DEUTETRABENAZINE
CONDYLOX, PODOFILOX
LOSEASONIQUE, ETHINYL ESTRADIOL
MICROZIDE, HYDROCHLOROTHIAZIDE
NOR-QD, NORETHINDRONE
NORETHINDRONE AND ETHINYL ESTRADIOL AND FERROUS FUMARATE, ETHINYL ESTRADIOL
PROAIR HFA, ALBUTEROL SULFATE
PROAIR RESPICLICK, ALBUTEROL SULFATE
PROGLYCEM, DIAZOXIDE
QNASL, BECLOMETHASONE DIPROPIONATE
QUARTETTE, ETHINYL ESTRADIOL
SEASONALE, ETHINYL ESTRADIOL
SEASONIQUE, ETHINYL ESTRADIOL
ZIAC, BISOPROLOL FUMARATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** T ******TEVA PARENTERAL**

* TEVA PARENTERAL MEDICINES INC
 LEVALBUTEROL HYDROCHLORIDE, LEVALBUTEROL HYDROCHLORIDE

TEVA PHARM

* TEVA PHARMACEUTICAL INDUSTRIES LTD
 AIRDUO RESPICLICK, FLUTICASONE PROPIONATE

TEVA PHARMS

* TEVA PHARMACEUTICALS DEVELOPMENT INC
 CHLORPROMAZINE HYDROCHLORIDE, CHLORPROMAZINE HYDROCHLORIDE
 DOXEPIN HYDROCHLORIDE, DOXEPIN HYDROCHLORIDE
 LISDEXAMFETAMINE DIMESYLATE, LISDEXAMFETAMINE DIMESYLATE
 PYRIMETHAMINE, PYRIMETHAMINE

* TEVA PHARMACEUTICALS USA
 AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE
 ANASTROZOLE, ANASTROZOLE
 BUDESONIDE, BUDESONIDE
 CARBAMAZEPINE, CARBAMAZEPINE
 CEFADROXIL, CEFADROXIL/CEFADROXIL HEMIHYDRATE
 CEFDINIR, CEFDINIR
 CROMOLYN SODIUM, CROMOLYN SODIUM
 FAMCICLOVIR, FAMCICLOVIR
 FLUVASTATIN SODIUM, FLUVASTATIN SODIUM
 GLIPIZIDE AND METFORMIN HYDROCHLORIDE, GLIPIZIDE
 HYDROXYCHLOROQUINE SULFATE, HYDROXYCHLOROQUINE SULFATE
 IRBESARTAN, IRBESARTAN
 LANSOPRAZOLE, LANSOPRAZOLE
 LETROZOLE, LETROZOLE
 LEVOCETIRIZINE DIHYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE
 LOSARTAN POTASSIUM AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
 MONTELUKAST SODIUM, MONTELUKAST SODIUM
 MYCOPHENOLATE MOFETIL, MYCOPHENOLATE MOFETIL
 OLANZAPINE AND FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 OXALIPLATIN, OXALIPLATIN
 PACLITAXEL, PACLITAXEL
 PRAZOSIN HYDROCHLORIDE, PRAZOSIN HYDROCHLORIDE
 QUETIAPINE FUMARATE, QUETIAPINE FUMARATE
 QUININE SULFATE, QUININE SULFATE
 RIZATRIPTAN BENZOATE, RIZATRIPTAN BENZOATE
 SILDENAFIL CITRATE, SILDENAFIL CITRATE
 TAMSULOSIN HYDROCHLORIDE, TAMSULOSIN HYDROCHLORIDE
 VANDAZOLE, METRONIDAZOLE
 VARDENAFIL HYDROCHLORIDE, VARDENAFIL HYDROCHLORIDE

TEVA PHARMS INC

* TEVA PHARMACEUTICALS INC
 ALVAIZ, ELTROMBOPAG CHOLINE
 ALYQ, TADALAFIL
 CETRORELIX ACETATE, CETRORELIX ACETATE
 MESALAMINE, MESALAMINE
 PACLITAXEL, PACLITAXEL
 PAZOPANIB HYDROCHLORIDE, PAZOPANIB HYDROCHLORIDE
 THEOPHYLLINE, THEOPHYLLINE
 TIOPRONIN, TIOPRONIN

TEVA PHARMS INTL

* TEVA PHARMACEUTICALS INTERNATIONAL GMBH
 AMRIX, CYCLOBENZAPRINE HYDROCHLORIDE

TEVA PHARMS USA

* TEVA PHARMACEUTICALS USA
 ACITRETIN, ACITRETIN
 ALMOTRIPTAN MALATE, ALMOTRIPTAN MALATE
 ARFORMOTEROL TARTRATE, ARFORMOTEROL TARTRATE
 ATAZANAVIR SULFATE, ATAZANAVIR SULFATE
 ATOMOXETINE HYDROCHLORIDE, ATOMOXETINE HYDROCHLORIDE
 ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM
 BEXAROTENE, BEXAROTENE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** T ****

- * TEVA PHARMACEUTICALS USA
 - BUDESONIDE, BUDESONIDE
 - CLARAVIS, ISOTRETINOIN
 - COPAXONE, GLATIRAMER ACETATE
 - DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE
 - DEXMETHYLPHENIDATE HYDROCHLORIDE, DEXMETHYLPHENIDATE HYDROCHLORIDE
 - EFAVIRENZ, EMTRICITABINE, AND TENOFOVIR DISOPROXIL FUMARATE, EFAVIRENZ
 - EPTIFIBATIDE, EPTIFIBATIDE
 - ESTRADIOL, ESTRADIOL
 - FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 - FLUVASTATIN SODIUM, FLUVASTATIN SODIUM
 - FOSAPREPITANT DIMEGLUMINE, FOSAPREPITANT DIMEGLUMINE
 - GUANFACINE HYDROCHLORIDE, GUANFACINE HYDROCHLORIDE
 - IBUPROFEN AND FAMOTIDINE, FAMOTIDINE
 - ICOSAPENT ETHYL, ICOSAPENT ETHYL
 - IMATINIB MESYLATE, IMATINIB MESYLATE
 - IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE
 - LANSOPRAZOLE, LANSOPRAZOLE
 - LEVALBUTEROL HYDROCHLORIDE, LEVALBUTEROL HYDROCHLORIDE
 - LOGILIA, ULIPRISTAL ACETATE
 - METOCLOPRAMIDE HYDROCHLORIDE, METOCLOPRAMIDE HYDROCHLORIDE
 - MOXIFLOXACIN HYDROCHLORIDE, MOXIFLOXACIN HYDROCHLORIDE
 - OLMESARTAN MEDOXOMIL, AMLODIPINE AND HYDROCHLOROTHIAZIDE, AMLODIPINE BESYLATE
 - OMEPRAZOLE, OMEPRAZOLE
 - PALONOSETRON HYDROCHLORIDE, PALONOSETRON HYDROCHLORIDE
 - PARICALCITOL, PARICALCITOL
 - PIOGLITAZONE HYDROCHLORIDE AND METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 - PIOGLITAZONE HYDROCHLORIDE, PIOGLITAZONE HYDROCHLORIDE
 - RALOXIFENE HYDROCHLORIDE, RALOXIFENE HYDROCHLORIDE
 - RISEDRONATE SODIUM, RISEDRONATE SODIUM
 - SILDENAFIL CITRATE, SILDENAFIL CITRATE
 - SOLIFENACIN SUCCINATE, SOLIFENACIN SUCCINATE
 - SULFAMETHOXAZOLE AND TRIMETHOPRIM, SULFAMETHOXAZOLE
 - SUNITINIB MALATE, SUNITINIB MALATE
 - TADALAFIL, TADALAFIL
 - TENOFOVIR DISOPROXIL FUMARATE, TENOFOVIR DISOPROXIL FUMARATE
 - TOBRAMYCIN, TOBRAMYCIN
 - TOLTERODINE TARTRATE, TOLTERODINE TARTRATE
 - TREPROSTINIL, TREPROSTINIL
 - VIGABATRIN, VIGABATRIN
 - VILAZODONE HYDROCHLORIDE, VILAZODONE HYDROCHLORIDE
 - ZANOSAR, STREPTOZOCIN
- * TEVA PHARMACEUTICALS USA INC
 - ABIRATERONE ACETATE, ABIRATERONE ACETATE
 - ALYQ, TADALAFIL
 - AMOXICILLIN AND CLAVULANATE POTASSIUM, AMOXICILLIN
 - CAPECITABINE, CAPECITABINE
 - DAPTOMYCIN, DAPTOMYCIN
 - DARUNAVIR, DARUNAVIR
 - DEFERASIROX, DEFERASIROX
 - DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE
 - EFINAACONAZOLE, EFINAACONAZOLE
 - ELETRIPTAN HYDROBROMIDE, ELETRIPTAN HYDROBROMIDE
 - EMTRICITABINE AND TENOFOVIR DISOPROXIL FUMARATE, EMTRICITABINE
 - EPINEPHRINE (AUTOINJECTOR), EPINEPHRINE
 - ESTRADIOL, ESTRADIOL
 - EVEROLIMUS, EVEROLIMUS
 - FINGOLIMOD HYDROCHLORIDE, FINGOLIMOD HYDROCHLORIDE
 - FLUTICASONE PROPIONATE AND SALMETEROL XINAFOATE, FLUTICASONE PROPIONATE
 - ICATIBANT ACETATE, ICATIBANT ACETATE
 - IVERMECTIN, IVERMECTIN
 - LIOTHYRONINE SODIUM, LIOTHYRONINE SODIUM
 - MESALAMINE, MESALAMINE
 - METHYLERGONOVINE MALEATE, METHYLERGONOVINE MALEATE
 - METRONIDAZOLE, METRONIDAZOLE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** T ****

* TEVA PHARMACEUTICALS USA INC
 MYCOPHENOLATE MOFETIL, MYCOPHENOLATE MOFETIL
 NALOXONE HYDROCHLORIDE, NALOXONE HYDROCHLORIDE (OTC)
 ONSURA, ETHINYL ESTRADIOL
 OSELTAMIVIR PHOSPHATE, OSELTAMIVIR PHOSPHATE
 OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE
 PALIPERIDONE PALMITATE, PALIPERIDONE PALMITATE
 PENCICLOVIR, PENCICLOVIR
 PIRFENIDONE, PIRFENIDONE
 TERIFLUNOMIDE, TERIFLUNOMIDE
 TERIPARATIDE, TERIPARATIDE
 TOBRAMYCIN, TOBRAMYCIN
 TRAZODONE HYDROCHLORIDE, TRAZODONE HYDROCHLORIDE
 TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE

TEVA PHARMS USA INC

* TEVA PHARMACEUTICALS USA INC
 CYCLOSPORINE, CYCLOSPORINE
 ERLOTINIB HYDROCHLORIDE, ERLOTINIB HYDROCHLORIDE
 ERYTHROMYCIN, ERYTHROMYCIN
 ETHINYL ESTRADIOL; ETONOGESTREL, ETHINYL ESTRADIOL
 FINZALA, ETHINYL ESTRADIOL
 FORMOTEROL FUMARATE, FORMOTEROL FUMARATE
 LAPATINIB DITOSYLATE, LAPATINIB DITOSYLATE
 LUBIPROSTONE, LUBIPROSTONE
 MIFEPRISTONE, MIFEPRISTONE
 OCTREOTIDE ACETATE, OCTREOTIDE ACETATE
 PLERIXAFOR, PLERIXAFOR
 POTASSIUM CITRATE, POTASSIUM CITRATE
 RISPERIDONE, RISPERIDONE
 SORAFENIB TOSYLATE, SORAFENIB TOSYLATE
 TASIMELTEON, TASIMELTEON
 TIOPRONIN, TIOPRONIN

TEYRO LABS

* TEYRO LABS PRIVATE LTD
 CAPECITABINE, CAPECITABINE
 CARBOPLATIN, CARBOPLATIN
 GEMCITABINE HYDROCHLORIDE, GEMCITABINE HYDROCHLORIDE
 TOPOTECAN HYDROCHLORIDE, TOPOTECAN HYDROCHLORIDE

THE FEINSTEIN INST

* THE FEINSTEIN INSTITUTE FOR MEDICAL RESEARCH
 SODIUM FLUORIDE F-18, SODIUM FLUORIDE F-18

THE J MOLNER

* THE J MOLNER CO OU
 DESOXIMETASONE, DESOXIMETASONE
 DIFLORASONE DIACETATE, DIFLORASONE DIACETATE
 HYDROCORTISONE BUTYRATE, HYDROCORTISONE BUTYRATE
 LIDOCAINE HYDROCHLORIDE, LIDOCAINE HYDROCHLORIDE

THEA PHARMA

* THEA PHARMA INC
 AKTEN, LIDOCAINE HYDROCHLORIDE
 AZASITE, AZITHROMYCIN
 BETIMOL, TIMOLOL
 COSOPT PF, DORZOLAMIDE HYDROCHLORIDE
 COSOPT, DORZOLAMIDE HYDROCHLORIDE
 IYUZEH, LATANOPROST
 ZIOPTAN, TAFLUPROST

THEPHARMANETWORK LLC

* THEPHARMANETWORK LLC
 BENZONATATE, BENZONATATE
 ISONIAZID, ISONIAZID
 METHADONE HYDROCHLORIDE, METHADONE HYDROCHLORIDE
 NIMODIPINE, NIMODIPINE
 THERMAZENE, SILVER SULFADIAZINE

THERACOSBIO

* THERACOSBIO LLC

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** T ****

* THERACOSBIO LLC

BRENZAVVY, BEXAGLIFLOZIN

THERAGNOSTICS

* THERAGNOSTICS INC

NEPHROSCAN, TECHNETIUM TC-99M SUCCIMER

THERAKOS DEVELOPMENT

* THERAKOS DEVELOPMENT LTD

UVADEX, METHOXSALEN

THERAVIA

* THERAVIA PHARMA

SIKLOS, HYDROXYUREA

THINQ PHARM-CRO PVT

* THINQ PHARMA-CRO PRIVATE LTD

CHLORPROMAZINE HYDROCHLORIDE, CHLORPROMAZINE HYDROCHLORIDE

CLOTRIMAZOLE, CLOTRIMAZOLE

FLUCONAZOLE, FLUCONAZOLE

MIDODRINE HYDROCHLORIDE, MIDODRINE HYDROCHLORIDE

TINIDAZOLE, TINIDAZOLE

TIANJIN KINGYORK

* TIANJIN KINGYORK PHARMACEUTICALS CO LTD

METHYLPREDNISOLONE SODIUM SUCCINATE, METHYLPREDNISOLONE SODIUM SUCCINATE

TIANJIN TIANYAO

* TIANJIN TIANYAO PHARMACEUTICALS CO LTD

CELECOXIB, CELECOXIB

METHYLPREDNISOLONE, METHYLPREDNISOLONE

TILDE SCIENCES

* TILDE SCIENCES LLC

DARAPRIM, PYRIMETHAMINE

TOLMAR

* TOLMAR INC

ELIGARD KIT, LEUPROLIDE ACETATE

JATENZO, TESTOSTERONE UNDECANOATE

* TOLMAR INTERNATIONAL LTD

FENSOLVI KIT, LEUPROLIDE ACETATE

TONIX MEDS

* TONIX MEDICINES INC

TOSYMRA, SUMATRIPTAN

ZEMBRACE SYMTOUCH, SUMATRIPTAN SUCCINATE

TOPROL

* TOPROL ACQUISITION LLC

TOPROL-XL, METOPROLOL SUCCINATE

TORRENT

* TORRENT PHARMA INC

APREPITANT, APREPITANT

ERYTHROMYCIN, ERYTHROMYCIN

ORMALVI, DICHLORPHENAMIDE

* TORRENT PHARMACEUTICALS LTD

ACYCLOVIR, ACYCLOVIR

ANAGRELIDE HYDROCHLORIDE, ANAGRELIDE HYDROCHLORIDE

APIXABAN, APIXABAN

ARIPIPIRAZOLE, ARIPIPIRAZOLE

CELECOXIB, CELECOXIB

CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE

DAPSONE, DAPSONE

DARIFENACIN HYDROBROMIDE, DARIFENACIN HYDROBROMIDE

DEFERASIROX, DEFERASIROX

DIMETHYL FUMARATE, DIMETHYL FUMARATE

DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE

FENOFIBRATE (MICRONIZED), FENOFIBRATE

FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE

ITRACONAZOLE, ITRACONAZOLE

LAMOTRIGINE, LAMOTRIGINE

LURASIDONE HYDROCHLORIDE, LURASIDONE HYDROCHLORIDE

MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** T ****

* TORRENT PHARMACEUTICALS LTD
 MOMETASONE FUROATE, MOMETASONE FUROATE
 MONTELUKAST SODIUM, MONTELUKAST SODIUM
 NEBIVOLOL HYDROCHLORIDE, NEBIVOLOL HYDROCHLORIDE
 NYSTATIN, NYSTATIN
 OLANZAPINE, OLANZAPINE
 OLMESARTAN MEDOXOMIL, AMLODIPINE AND HYDROCHLOROTHIAZIDE, AMLODIPINE BESYLATE
 PENCICLOVIR, PENCICLOVIR
 RABEPRAZOLE SODIUM, RABEPRAZOLE SODIUM
 ROFLUMILAST, ROFLUMILAST
 ROSUVASTATIN CALCIUM, ROSUVASTATIN CALCIUM
 SACUBITRIL AND VALSARTAN, SACUBITRIL
 SILDENAFIL CITRATE, SILDENAFIL CITRATE
 SORAFENIB TOSYLATE, SORAFENIB TOSYLATE
 TADALAFIL, TADALAFIL
 TIOPRONIN, TIOPRONIN
 TOLTERODINE TARTRATE, TOLTERODINE TARTRATE
 TRAZODONE HYDROCHLORIDE, TRAZODONE HYDROCHLORIDE

TORRENT PHARMS

* TORRENT PHARMACEUTICALS LTD
 CARBAMAZEPINE, CARBAMAZEPINE
 CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE
 DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
 ISOSORBIDE MONONITRATE, ISOSORBIDE MONONITRATE
 LAMOTRIGINE, LAMOTRIGINE
 LEVETIRACETAM, LEVETIRACETAM
 PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
 PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE
 RISPERIDONE, RISPERIDONE
 ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE

TORRENT PHARMS LTD

* TORRENT PHARMACEUTICALS LTD
 CLOPIDOGREL BISULFATE, CLOPIDOGREL BISULFATE
 ESCITALOPRAM OXALATE, ESCITALOPRAM OXALATE
 FELODIPINE, FELODIPINE
 MONTELUKAST SODIUM, MONTELUKAST SODIUM
 SILDENAFIL CITRATE, SILDENAFIL CITRATE

TRAVERE

* TRAVERE THERAPEUTICS INC
 FILSPARI, SPARSENTAN

TREVENA

* TREVENA INC
 OLINVYK, OLICERIDINE

TRIS PHARMA INC

* TRIS PHARMA INC
 DEFLAZACORT, DEFLAZACORT
 DEXMETHYLPHENIDATE HYDROCHLORIDE, DEXMETHYLPHENIDATE HYDROCHLORIDE
 DEXTROAMPHETAMINE SULFATE, DEXTROAMPHETAMINE SULFATE
 DEXTROMETHORPHAN POLISTIREX, DEXTROMETHORPHAN POLISTIREX (OTC)
 DYANAVEL XR 10, AMPHETAMINE
 DYANAVEL XR 15, AMPHETAMINE
 DYANAVEL XR 20, AMPHETAMINE
 DYANAVEL XR 5, AMPHETAMINE
 DYANAVEL XR, AMPHETAMINE
 HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
 HYDROCODONE POLISTIREX AND CHLORPHENIRAMINE POLISTIREX, CHLORPHENIRAMINE POLISTIREX
 IBUPROFEN, IBUPROFEN (OTC)
 METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
 MORPHINE SULFATE, MORPHINE SULFATE
 ONYDA XR, CLONIDINE HYDROCHLORIDE
 PROMETHAZINE HYDROCHLORIDE AND CODEINE PHOSPHATE, CODEINE PHOSPHATE
 PROMETHAZINE HYDROCHLORIDE AND DEXTROMETHORPHAN HYDROBROMIDE, DEXTROMETHORPHAN
 PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
 RISPERIDONE, RISPERIDONE
 THEOPHYLLINE, THEOPHYLLINE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** T ******TRUPHARMA**

* TRUPHARMA LLC
 AMINOCAPROIC ACID, AMINOCAPROIC ACID
 APOMORPHINE HYDROCHLORIDE, APOMORPHINE HYDROCHLORIDE
 BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
 BISOPROLOL FUMARATE, BISOPROLOL FUMARATE
 CLOMIPRAMINE HYDROCHLORIDE, CLOMIPRAMINE HYDROCHLORIDE
 CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
 CLOTRIMAZOLE, CLOTRIMAZOLE
 DAPSONE, DAPSONE
 FLUOCINOLONE ACETONIDE, FLUOCINOLONE ACETONIDE
 KETOCONAZOLE, KETOCONAZOLE
 METHYLERGONOVINE MALEATE, METHYLERGONOVINE MALEATE
 NAPROXEN, NAPROXEN
 OXAZEPAM, OXAZEPAM
 OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE

TRUSTEES UNIV PA

* TRUSTEES OF THE UNIV OF PENNSYLVANIA
 FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18

TWI PHARMS

* TWI PHARMACEUTICALS INC
 ATENOLOL AND CHLORTHALIDONE, ATENOLOL
 ATENOLOL, ATENOLOL
 BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
 DEXLANSOPRAZOLE, DEXLANSOPRAZOLE
 DICYCLOMINE HYDROCHLORIDE, DICYCLOMINE HYDROCHLORIDE
 DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
 DIMETHYL FUMARATE, DIMETHYL FUMARATE
 DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
 FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 FLUPHENAZINE HYDROCHLORIDE, FLUPHENAZINE HYDROCHLORIDE
 FORFIVO XL, BUPROPION HYDROCHLORIDE
 GUANFACINE HYDROCHLORIDE, GUANFACINE HYDROCHLORIDE
 MEGESTROL ACETATE, MEGESTROL ACETATE
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 MYCOPHENOLIC SODIUM, MYCOPHENOLIC SODIUM
 NAPRELAN, NAPROXEN SODIUM
 NIFEDIPINE, NIFEDIPINE
 POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
 TENORETIC 100, ATENOLOL
 TENORETIC 50, ATENOLOL
 TENORMIN, ATENOLOL
 TERBUTALINE SULFATE, TERBUTALINE SULFATE
 TESTOSTERONE, TESTOSTERONE
 TOPIRAMATE, TOPIRAMATE
 ZESTRIL, LISINOPRIL

TWI PHARMS INC

* TWI PHARMACEUTICALS INC
 CYCLOBENZAPRINE HYDROCHLORIDE, CYCLOBENZAPRINE HYDROCHLORIDE

**** U ******UBI**

* UBI PHARMA INC
 DESMOPRESSIN ACETATE, DESMOPRESSIN ACETATE
 LEUPROLIDE ACETATE, LEUPROLIDE ACETATE
 VORICONAZOLE, VORICONAZOLE

UCB INC

* UCB INC
 BRIVIACT, BRIVARACETAM
 FINTEPLA, FENFLURAMINE HYDROCHLORIDE
 KEPPRA XR, LEVETIRACETAM
 KEPPRA, LEVETIRACETAM
 NAYZILAM, MIDAZOLAM
 NEUPRO, ROTIGOTINE
 VIMPAT, LACOSAMIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** U ****

* UCB INC

ZILBRYSQ, ZILUCOPLAN SODIUM

UCLA BIOMEDICAL

* UCLA BIOMEDICAL CYCLOTRON

AMMONIA N 13, AMMONIA N-13

FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18

UCSF RODIOPHARM

* UCSF RODIOPHARMACEUTICAL FACILITY

AMMONIA N 13, AMMONIA N-13

FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18

UIHC PET IMAGING

* UNIV IOWA HOSPS AND CLINICS PET IMAGING CENTER

FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18

GALLIUM GA 68 EDOTREOTIDE, GALLIUM GA-68 EDOTREOTIDE

ULTRAGENYX PHARM INC

* ULTRAGENYX PHARMACEUTICAL INC

DOJOLVI, TRIHEPTANOIN

UMEDICA

* UMEDICA LABORATORIES PRIVATE LTD

ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM

CARBAMAZEPINE, CARBAMAZEPINE

CELECOXIB, CELECOXIB

CHLORTHALIDONE, CHLORTHALIDONE

DICLOFENAC POTASSIUM, DICLOFENAC POTASSIUM

EFINACONAZOLE, EFINACONAZOLE

GLYCOPYRROLATE, GLYCOPYRROLATE

NEOSTIGMINE METHYLSULFATE, NEOSTIGMINE METHYLSULFATE

OLMESARTAN MEDOXOMIL AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE

OLMESARTAN MEDOXOMIL, OLMESARTAN MEDOXOMIL

ROSUVASTATIN CALCIUM, ROSUVASTATIN CALCIUM

SILDENAFIL CITRATE, SILDENAFIL CITRATE

SUCCINYLCHOLINE CHLORIDE, SUCCINYLCHOLINE CHLORIDE

TADALAFIL, TADALAFIL

UNICHEM

* UNICHEM LABORATORIES LTD

ALFUZOSIN HYDROCHLORIDE, ALFUZOSIN HYDROCHLORIDE

ALLOPURINOL, ALLOPURINOL

AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE

AMITRIPTYLINE HYDROCHLORIDE, AMITRIPTYLINE HYDROCHLORIDE

AMLODIPINE BESYLATE, AMLODIPINE BESYLATE

ARIPIPIRAZOLE, ARIPIPIRAZOLE

ATENOLOL AND CHLORTHALIDONE, ATENOLOL

ATENOLOL, ATENOLOL

BACLOFEN, BACLOFEN

BISOPROLOL FUMARATE AND HYDROCHLOROTHIAZIDE, BISOPROLOL FUMARATE

BISOPROLOL FUMARATE, BISOPROLOL FUMARATE

BUSPIRONE HYDROCHLORIDE, BUSPIRONE HYDROCHLORIDE

CARBAMAZEPINE, CARBAMAZEPINE

CETIRIZINE HYDROCHLORIDE AND PSEUDOEPHEDRINE HYDROCHLORIDE, CETIRIZINE HYDROCHLORIDE

CHLORTHALIDONE, CHLORTHALIDONE

CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE

CYCLOBENZAPRINE HYDROCHLORIDE, CYCLOBENZAPRINE HYDROCHLORIDE

DIPHENOXYLATE HYDROCHLORIDE AND ATROPINE SULFATE, ATROPINE SULFATE

DIVALPROEX SODIUM, DIVALPROEX SODIUM

DOXAZOSIN MESYLATE, DOXAZOSIN MESYLATE

ETODOLAC, ETODOLAC

FAMOTIDINE, FAMOTIDINE

GUANFACINE HYDROCHLORIDE, GUANFACINE HYDROCHLORIDE

HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE

IRBESARTAN, IRBESARTAN

KETOTIFEN FUMARATE, KETOTIFEN FUMARATE (OTC)

LACOSAMIDE, LACOSAMIDE

LOSARTAN POTASSIUM AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE

LOSARTAN POTASSIUM, LOSARTAN POTASSIUM

MELOXICAM, MELOXICAM

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** U ****

* UNICHEM LABORATORIES LTD
 MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 METHSCOPOLAMINE BROMIDE, METHSCOPOLAMINE BROMIDE
 METOLAZONE, METOLAZONE
 METRONIDAZOLE, METRONIDAZOLE
 MONTELUKAST SODIUM, MONTELUKAST SODIUM
 NEBIVOLOL HYDROCHLORIDE, NEBIVOLOL HYDROCHLORIDE
 ORPHENADRINE CITRATE, ORPHENADRINE CITRATE
 PINDOLOL, PINDOLOL
 PIROXICAM, PIROXICAM
 PRASUGREL HYDROCHLORIDE, PRASUGREL HYDROCHLORIDE
 PRIMAQUINE PHOSPHATE, PRIMAQUINE PHOSPHATE
 QUETIAPINE FUMARATE, QUETIAPINE FUMARATE
 RANOLAZINE, RANOLAZINE
 SOLIFENACIN SUCCINATE, SOLIFENACIN SUCCINATE
 TADALAFIL, TADALAFIL
 TOLTERODINE TARTRATE, TOLTERODINE TARTRATE
 TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE
 ZALEPLON, ZALEPLON
 ZONISAMIDE, ZONISAMIDE

UNICHEM LABS LTD

* UNICHEM LABORATORIES LIMITED
 DIVALPROEX SODIUM, DIVALPROEX SODIUM
 * UNICHEM LABORATORIES LTD
 LAMOTRIGINE, LAMOTRIGINE
 TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE
 TOPIRAMATE, TOPIRAMATE

UNIMARK REMEDIES LTD

* UNIMARK REMEDIES LTD
 MONTELUKAST SODIUM, MONTELUKAST SODIUM

UNIQUE

* UNIQUE PHARMACEUTICAL LABORATORIES A DIV OF JB CHEMICALS AND PHARMACEUTICALS LTD
 ATENOLOL, ATENOLOL
 CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 CETIRIZINE HYDROCHLORIDE HIVES, CETIRIZINE HYDROCHLORIDE (OTC)
 CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE
 CLOMIPRAMINE HYDROCHLORIDE, CLOMIPRAMINE HYDROCHLORIDE
 DICLOFENAC SODIUM, DICLOFENAC SODIUM
 FEXOFENADINE HYDROCHLORIDE ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
 GLIPIZIDE, GLIPIZIDE
 LITHIUM CARBONATE, LITHIUM CARBONATE
 OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE
 TOLTERODINE TARTRATE, TOLTERODINE TARTRATE
 VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE

UNIQUE PHARM

* UNIQUE PHARMACEUTICAL LABORATORIES
 CARBAMAZEPINE, CARBAMAZEPINE
 DOXEPIN HYDROCHLORIDE, DOXEPIN HYDROCHLORIDE
 ENALAPRIL MALEATE, ENALAPRIL MALEATE
 LORATADINE, LORATADINE (OTC)

UNITED GUARDIAN

* UNITED GUARDIAN INC
 RENACIDIN, CITRIC ACID

UNITED THERAP

* UNITED THERAPEUTICS CORP
 ORENITRAM, TREPROSTINIL DIOLAMINE
 REMODULIN, TREPROSTINIL
 TYVASO DPI, TREPROSTINIL
 TYVASO, TREPROSTINIL

UNIV ALABAMA BIRM

* UNIV ALABAMA AT BIRMINGHAM
 AMMONIA N 13, AMMONIA N-13

UNIV CA LOS ANGELES

* UNIV CALIFORNIA LOS ANGELES

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** U ****

* UNIV CALIFORNIA LOS ANGELES

GALLIUM GA 68 GOZETOTIDE, GALLIUM GA-68 GOZETOTIDE

UNIV MICHIGAN

* UNIV MICHIGAN PET RADIOPHARMACEUTICAL PRODUCTION PROGRAM

FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18

UNIV OF CA SAN FRAN

* UNIV OF CALIFORNIA SAN FRANCISCO

GALLIUM GA 68 GOZETOTIDE, GALLIUM GA-68 GOZETOTIDE

UNIV SOUTHERN CA

* UNIV SOUTHERN CALIFORNIA DBA USC MOLECULAR IMAGING CENTER

FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18

UNIV TX MD ANDERSON

* UNIV TEXAS MD ANDERSON CANCER CENTER

FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18

UNIV TX SW MEDCTR

* UNIV TEXAS SOUTHWESTERN MEDCTR

AMMONIA N 13, AMMONIA N-13

UNIV UTAH CYCLOTRON

* UNIV UTAH CYCLOTRON RADIOCHEMISTRY LAB

FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18

SODIUM FLUORIDE F-18, SODIUM FLUORIDE F-18

UNIV WISCONSIN

* UNIV WISCONSIN SYSTEM

AMMONIA N 13, AMMONIA N-13

UPJOHN

* UPJOHN MANUFACTURING IRELAND UNLTD

LIPITOR, ATORVASTATIN CALCIUM

RELPAX, ELETRIPTAN HYDROBROMIDE

* UPJOHN US 2 LLC

CELEBREX, CELECOXIB

DETROL LA, TOLTERODINE TARTRATE

DETROL, TOLTERODINE TARTRATE

EFFEXOR XR, VENLAFAXINE HYDROCHLORIDE

INSPIRA, EPLERENONE

LYRICA CR, PREGABALIN

LYRICA, PREGABALIN

XALATAN, LATANOPROST

XANAX XR, ALPRAZOLAM

XANAX, ALPRAZOLAM

UPSHER SMITH LABS

* UPSHER SMITH LABORATORIES LLC

AMANTADINE HYDROCHLORIDE, AMANTADINE HYDROCHLORIDE

AZELASTINE HYDROCHLORIDE, AZELASTINE HYDROCHLORIDE

BACLOFEN, BACLOFEN

BETHANECHOL CHLORIDE, BETHANECHOL CHLORIDE

BEXAROTENE, BEXAROTENE

BRIMONIDINE TARTRATE AND TIMOLOL MALEATE, BRIMONIDINE TARTRATE

BUMETANIDE, BUMETANIDE

CHLORPROMAZINE HYDROCHLORIDE, CHLORPROMAZINE HYDROCHLORIDE

CHLORZOXAZONE, CHLORZOXAZONE

CLOBAZAM, CLOBAZAM

DEFLAZACORT, DEFLAZACORT

DIVALPROEX SODIUM, DIVALPROEX SODIUM

DOXAZOSIN MESYLATE, DOXAZOSIN MESYLATE

ELIGLUSTAT TARTRATE, ELIGLUSTAT TARTRATE

ETHACRYNIC ACID, ETHACRYNIC ACID

EXEMESTANE, EXEMESTANE

FAMOTIDINE, FAMOTIDINE

FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE

FLUPHENAZINE HYDROCHLORIDE, FLUPHENAZINE HYDROCHLORIDE

FLUVOXAMINE MALEATE, FLUVOXAMINE MALEATE

HALOPERIDOL, HALOPERIDOL

ISOTRETINOIN, ISOTRETINOIN

JANTOVEN, WARFARIN SODIUM

KLOR-CON M10, POTASSIUM CHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** U ****

* UPSHER SMITH LABORATORIES LLC
 Klor-con M15, POTASSIUM CHLORIDE
 Klor-con M20, POTASSIUM CHLORIDE
 Klor-con, POTASSIUM CHLORIDE
 LAMIVUDINE, LAMIVUDINE
 MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
 MORPHINE SULFATE, MORPHINE SULFATE
 MOXIFLOXACIN HYDROCHLORIDE, MOXIFLOXACIN HYDROCHLORIDE
 MYORISAN, ISOTRETINOIN
 NYSTATIN, NYSTATIN
 ORVATEN, MIDODRINE HYDROCHLORIDE
 OXYBUTYNYN CHLORIDE, OXYBUTYNYN CHLORIDE
 PACERONE, AMIODARONE HYDROCHLORIDE
 PREVALITE, CHOLESTYRAMINE
 PROPAFENONE HYDROCHLORIDE, PROPAFENONE HYDROCHLORIDE
 QUDEXY XR, TOPIRAMATE
 RAMELTEON, RAMELTEON
 SODIUM POLYSTYRENE SULFONATE, SODIUM POLYSTYRENE SULFONATE
 VIGABATRIN, VIGABATRIN
 VOGELXO, TESTOSTERONE

UROGEN PHARMA

* UROGEN PHARMA LTD
 JELMYTO, MITOMYCIN

UROVANT

* UROVANT SCIENCES GMBH
 GEMTESA, VIBEGRON

US ANTIBIOTICS

* US ANTIBIOTICS LLC
 AMOXIL, AMOXICILLIN
 AUGMENTIN '125', AMOXICILLIN
 AUGMENTIN '250', AMOXICILLIN
 AUGMENTIN '875', AMOXICILLIN
 AUGMENTIN ES-600, AMOXICILLIN
 LAROTID, AMOXICILLIN

USPHARMA

* USPHARMA LTD
 NITRO-DUR, NITROGLYCERIN

USV

* USV PRIVATE LTD
 OLOPATADINE HYDROCHLORIDE, OLOPATADINE HYDROCHLORIDE (OTC)
 ZOLEDRONIC ACID, ZOLEDRONIC ACID

USWM

* USWM LLC
 IWILFIN, EFLORNITHINE HYDROCHLORIDE
 LUCEMYRA, LOFEXIDINE HYDROCHLORIDE
 REVONTO, DANTROLENE SODIUM

UTILITY THERAP

* UTILITY THERAPEUTICS LTD
 PIVYA, PIVMECILLINAM HYDROCHLORIDE

UTOPIC PHARMS

* UTOPIA PHARMACEUTICALS INC
 DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
 TOLTERODINE TARTRATE, TOLTERODINE TARTRATE
 TROSPIMUM CHLORIDE, TROSPIMUM CHLORIDE

**** V ******VALEANT**

* VALEANT PHARMACEUTICALS INTERNATIONAL
 BONTRIL PDM, PHENDIMETRAZINE TARTRATE
 MYSOLINE, PRIMIDONE

VALEANT BERMUDA

* VALEANT INTERNATIONAL BERMUDA
 RETIN-A, TRETINOIN

VALEANT INTL

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** V ****

* VALEANT INTERNATIONAL BARBADOS SRL
RETIN-A, TRETINOIN

* VALEANT INTERNATIONAL SRL
BENZAMYCIN, BENZOYL PEROXIDE

VALEANT LUXEMBOURG

* VALEANT PHARMACEUTICALS LUXEMBOURG SARL
TARGRETIN, BEXAROTENE

VALEANT PHARM INTL

* VALEANT PHARMACEUTICALS INTERNATIONAL
ANDROID 25, METHYLTESTOSTERONE
LIBRIUM, CHLORDIAZEPOXIDE HYDROCHLORIDE

VALEANT PHARMS

* VALEANT PHARMACEUTICALS NORTH AMERICA LLC
PENTOXIFYLLINE, PENTOXIFYLLINE

VALEANT PHARMS INTL

* VALEANT PHARMACEUTICALS INTERNATIONAL
COLAZAL, BALSALAZIDE DISODIUM
CUPRIMINE, PENICILLAMINE

VALEANT PHARMS NORTH

* VALEANT PHARMACEUTICALS NORTH AMERICA LLC
DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
FENOFIBRATE, FENOFIBRATE
NIFEDIPINE, NIFEDIPINE
RENOVA, TRETINOIN
RETIN-A, TRETINOIN

VALIDUS PHARMS

* VALIDUS PHARMACEUTICALS LLC
BUMEX, BUMETANIDE
EQUETRO, CARBAMAZEPINE
LASIX, FUROSEMIDE
LOPRESSOR HCT, HYDROCHLOROTHIAZIDE
LOPRESSOR, METOPROLOL TARTRATE
LOTENSIN HCT, BENAZEPRIL HYDROCHLORIDE
LOTENSIN, BENAZEPRIL HYDROCHLORIDE
NORPRAMIN, DESIPRAMINE HYDROCHLORIDE

VALIDUS PHARMS INC

* VALIDUS PHARMACEUTICALS INC
MARPLAN, ISOCARBOXAZID

VALINOR

* VALINOR PHARMA LLC
MOVANTIK, NALOXEGOL OXALATE

VANCOCIN ITALIA

* VANCOCIN ITALIA SRL
MULPLETA, LUSUTROMBOPAG

VANDA PHARMS INC

* VANDA PHARMACEUTICALS INC
FANAPT, ILOPERIDONE
HETLIOZ LQ, TASIMELTEON
HETLIOZ, TASIMELTEON
PONVORY, PONESIMOD

VELOXIS PHARMS INC

* VELOXIS PHARMACEUTICALS INC
ENVARUS XR, TACROLIMUS

VELZEN PHARMA PVT

* VELZEN PHARMA PVT LTD
DOXEPIN HYDROCHLORIDE, DOXEPIN HYDROCHLORIDE
GLYCOPYRROLATE, GLYCOPYRROLATE
NIFEDIPINE, NIFEDIPINE
THEOPHYLLINE, THEOPHYLLINE

VERITY

* VERITY PHARMACEUTICALS INC
TLANDO, TESTOSTERONE UNDECANOATE
TRELSTAR, TRIPTORELIN PAMOATE

VERO BIOTECH INC

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** V ****

* VERO BIOTECH INC
GENOSYL, NITRIC OXIDE

VERONA PHARMA

* VERONA PHARMA INC
OHTUVAYRE, ENSIFENTRINE

VEROSCIENCE

* VEROSCIENCE LLC
CYCLOSET, BROMOCRIPTINE MESYLATE

VERRICA PHARMS

* VERRICA PHARMACEUTICALS INC
YCANTH, CANTHARIDIN

VERTEX PHARMS

* VERTEX PHARMACEUTICALS INC
KALYDECO, IVACAFTOR

VERTEX PHARMS INC

* VERTEX PHARMACEUTICALS INC
ALYFTREK, DEUTIVACAFTOR
KALYDECO, IVACAFTOR
ORKAMBI, IVACAFTOR
SYMDEKO (COPACKAGED), IVACAFTOR
TRIKAFTA (COPACKAGED), ELEXACAFTOR, IVACAFTOR, TEZACAFTOR

VERTICAL PHARMS

* VERTICAL PHARMACEUTICALS LLC
DIVIGEL, ESTRADIOL

VIATRIS

* VIATRIS SPECIALTY LLC
CARDURA XL, DOXAZOSIN MESYLATE
CARDURA, DOXAZOSIN MESYLATE
DILANTIN, PHENYTOIN SODIUM
DILANTIN-125, PHENYTOIN
GEODON, ZIPRASIDONE HYDROCHLORIDE
GEODON, ZIPRASIDONE MESYLATE
NEURONTIN, GABAPENTIN
NITROSTAT, NITROGLYCERIN
NORVASC, AMLODIPINE BESYLATE
REVATIO, SILDENAFIL CITRATE
VIAGRA, SILDENAFIL CITRATE
ZOLOFT, SERTRALINE HYDROCHLORIDE

VICURON HOLDINGS

* VICURON HOLDINGS LLC
ERAXIS, ANIDULAFUNGIN

VIFOR FRESENIUS

* VIFOR FRESENIUS MEDICAL CARE RENAL PHARMA FRANCE
VELPHORO, FERRIC OXYHYDROXIDE

VIFOR PHARMA

* VIFOR PHARMA INC
VELTASSA, PATIROMER SORBITEX CALCIUM

VIIV HLTHCARE

* VIIV HEALTHCARE CO
APRETUDE, CABOTEGRAVIR
CABENUVA KIT, CABOTEGRAVIR
DOVATO, DOLUTEGRAVIR SODIUM
EPIVIR, LAMIVUDINE
JULUCA, DOLUTEGRAVIR SODIUM
RETROVIR, ZIDOVUDINE
RUKOBIA, FOSTEMSAVIR TROMETHAMINE
SELZENTRY, MARAVIROC
TIVICAY PD, DOLUTEGRAVIR SODIUM
TIVICAY, DOLUTEGRAVIR SODIUM
TRIUMEQ PD, ABACAVIR SULFATE
TRIUMEQ, ABACAVIR SULFATE
VOCABRIA, CABOTEGRAVIR SODIUM
ZIAGEN, ABACAVIR SULFATE

VIRTUS

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** V ****

* VIRTUS PHARMACEUTICALS LLC
 CROMOLYN SODIUM, CROMOLYN SODIUM
 LEVORPHANOL TARTRATE, LEVORPHANOL TARTRATE
 PHENDIMETRAZINE TARTRATE, PHENDIMETRAZINE TARTRATE
 PROMETRIUM, PROGESTERONE

VIRTUS PHARM

* VIRTUS PHARMACEUTICAL INC
 ALBUTEROL SULFATE, ALBUTEROL SULFATE

VISTA PHARMS

* VISTA PHARMACEUTICALS INC
 SULFAMETHOXAZOLE AND TRIMETHOPRIM, SULFAMETHOXAZOLE

VISTAPHARM

* VISTAPHARM INC
 ARIPIPIRAZOLE, ARIPIPIRAZOLE
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 METHADONE HYDROCHLORIDE, METHADONE HYDROCHLORIDE
 MYCOPHENOLATE MOFETIL, MYCOPHENOLATE MOFETIL

VISTAPHARM LLC

* VISTAPHARM LLC
 DIGOXIN, DIGOXIN
 METHADONE HYDROCHLORIDE, METHADONE HYDROCHLORIDE
 SUCRALFATE, SUCRALFATE

VISUM PHARM

* VISUM PHARMACEUTICAL CO LTD
 METOPROLOL SUCCINATE, METOPROLOL SUCCINATE

VITRUVIAS THERAP

* VITRUVIAS THERAPEUTICS LLC
 CYANOCOBALAMIN, CYANOCOBALAMIN

VIVUS LLC

* VIVUS LLC
 QSYMIA, PHENTERMINE HYDROCHLORIDE

VIWIT PHARM

* VIWIT PHARMACEUTICAL CO LTD
 LEVETIRACETAM, LEVETIRACETAM
 NITROGLYCERIN, NITROGLYCERIN
 PROCHLORPERAZINE EDISYLATE, PROCHLORPERAZINE EDISYLATE
 SERTRALINE HYDROCHLORIDE, SERTRALINE HYDROCHLORIDE
 TOPIRAMATE, TOPIRAMATE

VKT PHARMA

* VKT PHARMA PRIVATE LTD
 FAMOTIDINE, FAMOTIDINE (OTC)
 RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
 RANOLAZINE, RANOLAZINE
 SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
 TADALAFIL, TADALAFIL

VPNA

* VALEANT PHARMACEUTICALS NORTH AMERICA
 DICLOFENAC SODIUM, DICLOFENAC SODIUM

**** W ******WA UNIV SCH MED**

* WASHINGTON UNIV SCHOOL MEDICINE
 AMMONIA N 13, AMMONIA N-13

WANBANG BIOPHARMS

* WANBANG BIOPHARMACEUTICALS
 CARISOPRODOL, CARISOPRODOL
 LEFLUNOMIDE, LEFLUNOMIDE
 SUNITINIB MALATE, SUNITINIB MALATE

* WANBANG BIOPHARMACEUTICALS INC
 MYCOPHENOLIC SODIUM, MYCOPHENOLIC SODIUM

WATSON LABS

* WATSON LABORATORIES
 FOLIC ACID, FOLIC ACID
 PROPAFENONE HYDROCHLORIDE, PROPAFENONE HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** W ****

- * WATSON LABORATORIES INC
ACARBOSE, ACARBOSE
ALENDRONATE SODIUM, ALENDRONATE SODIUM
ALLOPURINOL, ALLOPURINOL
AMLODIPINE BESYLATE AND BENAZEPRIL HYDROCHLORIDE, AMLODIPINE BESYLATE
AMOXAPINE, AMOXAPINE
ATENOLOL AND CHLORTHALIDONE, ATENOLOL
CARISOPRODOL, CARISOPRODOL
CHLORZOXAZONE, CHLORZOXAZONE
CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE
COL-PROBENECID, COLCHICINE
DESOGESTREL AND ETHINYL ESTRADIOL, DESOGESTREL
DICYCLOMINE HYDROCHLORIDE, DICYCLOMINE HYDROCHLORIDE
DROSPIRENONE AND ETHINYL ESTRADIOL, DROSPIRENONE
ESTAZOLAM, ESTAZOLAM
GALANTAMINE HYDROBROMIDE, GALANTAMINE HYDROBROMIDE
GLIPIZIDE, GLIPIZIDE
GUANFACINE HYDROCHLORIDE, GUANFACINE HYDROCHLORIDE
LABETALOL HYDROCHLORIDE, LABETALOL HYDROCHLORIDE
LAMOTRIGINE, LAMOTRIGINE
LEVONORGESTREL AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
LISINOPRIL, LISINOPRIL
LORAZEPAM, LORAZEPAM
LOSARTAN POTASSIUM, LOSARTAN POTASSIUM
LOXAPINE SUCCINATE, LOXAPINE SUCCINATE
METHYLPREDNISOLONE, METHYLPREDNISOLONE
METRONIDAZOLE, METRONIDAZOLE
MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE
MINOXIDIL, MINOXIDIL
NABUMETONE, NABUMETONE
NALOXONE HYDROCHLORIDE AND PENTAZOCINE HYDROCHLORIDE, NALOXONE HYDROCHLORIDE
NATEGLINIDE, NATEGLINIDE
NEOMYCIN AND POLYMYXIN B SULFATE, NEOMYCIN SULFATE
NIZATIDINE, NIZATIDINE
NORETHINDRONE AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
ORPHENADRINE CITRATE, ORPHENADRINE CITRATE
PRAVASTATIN SODIUM, PRAVASTATIN SODIUM
PREDNISOLONE, PREDNISOLONE
PREDNISON, PREDNISON
PRIMIDONE, PRIMIDONE
PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
PROPRANOLOL HYDROCHLORIDE, PROPRANOLOL HYDROCHLORIDE
QUASENSE, ETHINYL ESTRADIOL
RAMIPRIL, RAMIPRIL
RIVASTIGMINE TARTRATE, RIVASTIGMINE TARTRATE
SULFASALAZINE, SULFASALAZINE
SULINDAC, SULINDAC
SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
TETRACYCLINE HYDROCHLORIDE, TETRACYCLINE HYDROCHLORIDE
TRIAMTERENE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
TRIHEXYPHENIDYL HYDROCHLORIDE, TRIHEXYPHENIDYL HYDROCHLORIDE
TRIMETHOPRIM, TRIMETHOPRIM
VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE
ZOVIA 1/50E-28, ETHINYL ESTRADIOL
- * WATSON LABS INC
LISINOPRIL AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE

WATSON LABS INC

- * WATSON LABORATORIES INC
AMANTADINE HYDROCHLORIDE, AMANTADINE HYDROCHLORIDE
AMBRISENTAN, AMBRISENTAN
AMLODIPINE BESYLATE AND BENAZEPRIL HYDROCHLORIDE, AMLODIPINE BESYLATE
BOSENTAN, BOSENTAN
BRINZOLAMIDE, BRINZOLAMIDE
BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
CELECOXIB, CELECOXIB

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** W ****

* WATSON LABORATORIES INC
 CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE
 COLCHICINE, COLCHICINE
 DICLOFENAC SODIUM, DICLOFENAC SODIUM
 DROSPIRENONE, ETHINYL ESTRADIOL AND LEVOMEFOLATE CALCIUM, DROSPIRENONE
 EZETIMIBE AND SIMVASTATIN, EZETIMIBE
 EZETIMIBE, EZETIMIBE
 METRONIDAZOLE, METRONIDAZOLE
 MODAFINIL, MODAFINIL
 NITROFURANTOIN (MONOHYDRATE/MACROCRYSTALS), NITROFURANTOIN
 PENICILLAMINE, PENICILLAMINE
 PERPHENAZINE, PERPHENAZINE
 PROPAFENONE HYDROCHLORIDE, PROPAFENONE HYDROCHLORIDE
 RALOXIFENE HYDROCHLORIDE, RALOXIFENE HYDROCHLORIDE
 ROSUVASTATIN CALCIUM, ROSUVASTATIN CALCIUM
 SILDENAFIL CITRATE, SILDENAFIL CITRATE
 SOLIFENACIN SUCCINATE, SOLIFENACIN SUCCINATE
 TADALAFIL, TADALAFIL

WATSON LABS TEVA

* WATSON LABORATORIES INC AN INDIRECT WHOLLY OWNED SUB OF TEVA PHARMACEUTICALS USA INC
 ALVIMOPAN, ALVIMOPAN
 BICALUTAMIDE, BICALUTAMIDE
 BUPRENORPHINE, BUPRENORPHINE
 CINACALCET HYDROCHLORIDE, CINACALCET HYDROCHLORIDE
 GLIPIZIDE, GLIPIZIDE
 ISRADIPINE, ISRADIPINE
 LEVOTHYROXINE SODIUM, LEVOTHYROXINE SODIUM **
 MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE
 NORETHINDRONE AND ETHINYL ESTRADIOL (10/11), ETHINYL ESTRADIOL
 NORETHINDRONE AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
 PROBENECID, PROBENECID
 TRIENTINE HYDROCHLORIDE, TRIENTINE HYDROCHLORIDE

WAYLIS THERAP

* WAYLIS THERAPEUTICS LLC
 ACIPHEX, RABEPRAZOLE SODIUM
 AVODART, DUTASTERIDE
 COREG, CARVEDILOL
 FLUTAMIDE, FLUTAMIDE
 JALYN, DUTASTERIDE
 LEUKERAN, CHLORAMBUCIL
 LUNESTA, ESZOPICLONE
 MYLERAN, BUSULFAN
 THIOGUANINE, THIOGUANINE
 VALIUM, DIAZEPAM

WES PHARMA INC

* WES PHARMA INC
 ACETAMINOPHEN AND CODEINE PHOSPHATE, ACETAMINOPHEN
 BROMPHENIRAMINE MALEATE, PSEUDOEPHEDRINE HYDROCHLORIDE AND DEXTROMETHORPHAN
 BUPRENORPHINE HYDROCHLORIDE AND NALOXONE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE
 HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
 METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
 OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN
 OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
 POTASSIUM CHLORIDE, POTASSIUM CHLORIDE

WEST WARD

* WEST WARD PHARMACEUTICAL CORP
 DICYCLOMINE HYDROCHLORIDE, DICYCLOMINE HYDROCHLORIDE

WEST WARD PHARM CORP

* WEST WARD PHARMACEUTICAL CORP
 ROCURONIUM BROMIDE, ROCURONIUM BROMIDE

WEST-WARD PHARMS INT

* WEST-WARD PHARMACEUTICALS INTERNATIONAL LTD
 BUMETANIDE, BUMETANIDE
 CYANOCOBALAMIN, CYANOCOBALAMIN
 CYTARABINE, CYTARABINE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** W ****

* WEST-WARD PHARMACEUTICALS INTERNATIONAL LTD
 DIPHENHYDRAMINE HYDROCHLORIDE, DIPHENHYDRAMINE HYDROCHLORIDE
 DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
 LIDOCAINE HYDROCHLORIDE, LIDOCAINE HYDROCHLORIDE
 MEPERIDINE HYDROCHLORIDE PRESERVATIVE FREE, MEPERIDINE HYDROCHLORIDE
 MEPERIDINE HYDROCHLORIDE, MEPERIDINE HYDROCHLORIDE
 OCTREOTIDE ACETATE (PRESERVATIVE FREE), OCTREOTIDE ACETATE
 OCTREOTIDE ACETATE, OCTREOTIDE ACETATE
 PENTOSTATIN, PENTOSTATIN
 THIAMINE HYDROCHLORIDE, THIAMINE HYDROCHLORIDE
 THIOTEPA, THIOTEPA

WESTMINSTER PHARMS

* WESTMINSTER PHARMACEUTICALS LLC
 EPLERENONE, EPLERENONE
 MIGLITOL, MIGLITOL
 POSACONAZOLE, POSACONAZOLE

WILSHIRE PHARMS INC

* WILSHIRE PHARMACEUTICALS INC
 ARFORMOTEROL TARTRATE, ARFORMOTEROL TARTRATE
 DEOXYCHOLIC ACID, DEOXYCHOLIC ACID
 DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE
 FORMOTEROL FUMARATE, FORMOTEROL FUMARATE
 ICATIBANT ACETATE, ICATIBANT ACETATE
 MERZEE, ETHINYL ESTRADIOL
 METHYLPREDNISOLONE ACETATE, METHYLPREDNISOLONE ACETATE
 OMEGA-3-ACID ETHYL ESTERS, OMEGA-3-ACID ETHYL ESTERS
 PERPHENAZINE, PERPHENAZINE
 TESTOSTERONE CYPIONATE, TESTOSTERONE CYPIONATE

WINDER LABS LLC

* WINDER LABORATOIRES LLC
 CHLORDIAZEPOXIDE HYDROCHLORIDE AND CLIDINIUM BROMIDE, CHLORDIAZEPOXIDE HYDROCHLORIDE
 DEXTROAMPHETAMINE SULFATE, DEXTROAMPHETAMINE SULFATE
 DIPHENOXYLATE HYDROCHLORIDE AND ATROPINE SULFATE, ATROPINE SULFATE
 MORPHINE SULFATE, MORPHINE SULFATE

WISCONSIN

* WISCONSIN MEDICAL RADIOPHARMACY LLC
 FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18

WOCKHARDT

* WOCKHARDT LTD
 CAPTOPRIL, CAPTOPRIL
 CEFTRIAZONE, CEFTRIAZONE SODIUM
 CHILDREN'S FEXOFENADINE HYDROCHLORIDE ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
 CHILDREN'S FEXOFENADINE HYDROCHLORIDE HIVES, FEXOFENADINE HYDROCHLORIDE (OTC)
 DIVALPROEX SODIUM, DIVALPROEX SODIUM
 FEXOFENADINE HYDROCHLORIDE ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
 FEXOFENADINE HYDROCHLORIDE HIVES, FEXOFENADINE HYDROCHLORIDE (OTC)
 FOSPHENYTOIN SODIUM, FOSPHENYTOIN SODIUM
 FUROSEMIDE, FUROSEMIDE
 KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
 LANSOPRAZOLE, LANSOPRAZOLE
 METOPROLOL SUCCINATE, METOPROLOL SUCCINATE
 NICARDIPINE HYDROCHLORIDE, NICARDIPINE HYDROCHLORIDE
 ONDANSETRON HYDROCHLORIDE PRESERVATIVE FREE, ONDANSETRON HYDROCHLORIDE
 ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
 SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE

WOCKHARDT BIO AG

* WOCKHARDT BIO AG
 ABIRATERONE ACETATE, ABIRATERONE ACETATE
 ACETAMINOPHEN, ACETAMINOPHEN
 BROMFED-DM, BROMPHENIRAMINE MALEATE
 CROMOLYN SODIUM, CROMOLYN SODIUM
 DECITABINE, DECITABINE
 ENTACAPONE, ENTACAPONE
 HYDROCODONE BITARTRATE AND HOMATROPINE METHYLBROMIDE, HOMATROPINE METHYLBROMIDE
 LIDOCAINE HYDROCHLORIDE, LIDOCAINE HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** W ****

* WOCKHARDT BIO AG
 LISINOPRIL, LISINOPRIL
 LOPERAMIDE HYDROCHLORIDE, LOPERAMIDE HYDROCHLORIDE (OTC)
 LORATADINE, LORATADINE (OTC)
 NYSTATIN, NYSTATIN
 OXACILLIN SODIUM, OXACILLIN SODIUM
 PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
 PIPERACILLIN AND TAZOBACTAM, PIPERACILLIN SODIUM
 PROMETHAZINE HYDROCHLORIDE AND CODEINE PHOSPHATE, CODEINE PHOSPHATE
 PROMETHAZINE PLAIN, PROMETHAZINE HYDROCHLORIDE
 PROMETHAZINE W/ DEXTROMETHORPHAN, DEXTROMETHORPHAN HYDROBROMIDE
 TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE
 VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE

WOCKHARDT LTD

* WOCKHARDT LTD
 BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
 ENALAPRIL MALEATE, ENALAPRIL MALEATE

WOODWARD

* WOODWARD PHARMA SERVICES LLC
 FLUCONAZOLE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, FLUCONAZOLE
 LOVAZA, OMEGA-3-ACID ETHYL ESTERS

WUSM CYCLOTRON

* WASHINGTON UNIV SCH MEDICINE CYCLOTRON FACILITY
 FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18

WUXI

* WUXI FORTUNE PHARMACEUTICAL CO LTD
 MYCOPHENOLATE MOFETIL, MYCOPHENOLATE MOFETIL

WYETH PHARMS

* WYETH PHARMACEUTICALS LLC
 DUAVEE, BAZEDOXIFENE ACETATE
 PREMARIN, ESTROGENS, CONJUGATED
 PREMPHASE 14/14, ESTROGENS, CONJUGATED
 PREMPRO, ESTROGENS, CONJUGATED
 PROTONIX IV, PANTOPRAZOLE SODIUM
 PROTONIX, PANTOPRAZOLE SODIUM
 TRECATOR, ETHIONAMIDE

**** X ******X-GEN PHARMS INC**

* X-GEN PHARMACEUTICALS INC
 PENTAMIDINE ISETHIONATE, PENTAMIDINE ISETHIONATE

X4 PHARMS

* X4 PHARMACEUTICALS INC
 XOLREMDI, MAVORIXAFOR

XCOVERY

* XCOVERY HOLDINGS INC
 ENSACOVE, ENSARTINIB HYDROCHLORIDE

XELLIA PHARMS APS

* XELLIA PHARMACEUTICALS APS
 COLISTIMETHATE SODIUM, COLISTIMETHATE SODIUM
 DAPTOMYCIN, DAPTOMYCIN
 MICAFUNGIN SODIUM, MICAFUNGIN SODIUM
 POLYMYXIN B SULFATE, POLYMYXIN B SULFATE
 TOBRAMYCIN SULFATE, TOBRAMYCIN SULFATE
 VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE

XERIS

* XERIS PHARMACEUTICALS INC
 GVOKE HYPOPEN, GLUCAGON
 GVOKE KIT, GLUCAGON
 GVOKE PFS, GLUCAGON
 KEVEYIS, DICHLORPHENAMIDE

XGEN PHARMS

* XGEN PHARMACEUTICALS DJB INC
 ACETAZOLAMIDE SODIUM, ACETAZOLAMIDE SODIUM

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** X ****

* XGEN PHARMACEUTICALS DJB INC
 AMPHOTERICIN B, AMPHOTERICIN B
 CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
 CYCLOPHOSPHAMIDE, CYCLOPHOSPHAMIDE
 DACTINOMYCIN, DACTINOMYCIN
 EDARAVONE, EDARAVONE
 FOLIC ACID, FOLIC ACID
 IBUPROFEN LYSINE, IBUPROFEN LYSINE
 LEVETIRACETAM, LEVETIRACETAM
 LINCOMYCIN HYDROCHLORIDE, LINCOMYCIN HYDROCHLORIDE
 LIOTHYRONINE SODIUM, LIOTHYRONINE SODIUM
 NELARABINE, NELARABINE
 NEOMYCIN AND POLYMYXIN B SULFATE, NEOMYCIN SULFATE
 PENTAMIDINE ISETHIONATE, PENTAMIDINE ISETHIONATE
 PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
 STREPTOMYCIN SULFATE, STREPTOMYCIN SULFATE
 TOBRAMYCIN SULFATE, TOBRAMYCIN SULFATE
 TRANEXAMIC ACID, TRANEXAMIC ACID

XIAMEN LP PHARM CO

* XIAMEN LP PHARMACEUTICAL CO LTD
 CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
 MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
 OPIPZA, ARIPIPRAZOLE
 PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE
 TOPIRAMATE, TOPIRAMATE

XIROMED

* XIROMED LLC
 DESOGESTREL AND ETHINYL ESTRADIOL, DESOGESTREL
 DROSPIRENONE AND ETHINYL ESTRADIOL, DROSPIRENONE
 ESTRADIOL AND NORETHINDRONE ACETATE, ESTRADIOL
 ETHYNODIOL DIACETATE AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
 LEVONORGESTREL AND ETHINYL ESTRADIOL AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
 LEVONORGESTREL AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
 LEVONORGESTREL, LEVONORGESTREL (OTC)
 MEDROXYPROGESTERONE ACETATE, MEDROXYPROGESTERONE ACETATE
 NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
 NORETHINDRONE ACETATE, NORETHINDRONE ACETATE
 NORETHINDRONE AND ETHINYL ESTRADIOL AND FERROUS FUMARATE, ETHINYL ESTRADIOL
 NORETHINDRONE, NORETHINDRONE

* XIROMED PHARMA ESPANA SL
 ACYCLOVIR, ACYCLOVIR
 ALTAVERA, ETHINYL ESTRADIOL
 BETAMETHASONE VALERATE, BETAMETHASONE VALERATE
 CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
 ENILLORING, ETHINYL ESTRADIOL
 EPHEDRINE SULFATE, EPHEDRINE SULFATE
 ESTARYLLA, ETHINYL ESTRADIOL
 ESTRADIOL VALERATE, ESTRADIOL VALERATE
 FLUOCINONIDE, FLUOCINONIDE
 FOSFOMYCIN TROMETHAMINE, FOSFOMYCIN TROMETHAMINE
 FULVESTRANT, FULVESTRANT
 GEMMILY, ETHINYL ESTRADIOL
 GLYCOPYRROLATE, GLYCOPYRROLATE
 GUANFACINE HYDROCHLORIDE, GUANFACINE HYDROCHLORIDE
 INTROVALE, ETHINYL ESTRADIOL
 ISIBLOOM, DESOGESTREL
 JAIMIESS, ETHINYL ESTRADIOL
 KETOCONAZOLE, KETOCONAZOLE
 LANSOPRAZOLE, LANSOPRAZOLE
 LEVONORGESTREL AND ETHINYL ESTRADIOL AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
 LEVONORGESTREL AND ETHINYL ESTRADIOL AND FERROUS FUMARATE, ETHINYL ESTRADIOL
 LEVONORGESTREL, LEVONORGESTREL (OTC)
 LEVOTHYROXINE SODIUM, LEVOTHYROXINE SODIUM
 LORYNA, DROSPIRENONE
 MIDODRINE HYDROCHLORIDE, MIDODRINE HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** X ****

* XIROMED PHARMA ESPANA SL
 NAFTIFINE HYDROCHLORIDE, NAFTIFINE HYDROCHLORIDE
 NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL AND FERROUS FUMARATE, ETHINYL ESTRADIOL
 OMEPRAZOLE, OMEPRAZOLE
 PROGESTERONE, PROGESTERONE
 RAMELTEON, RAMELTEON
 SODIUM NITROPRUSSIDE, SODIUM NITROPRUSSIDE
 SYEDA, DROSPIRENONE
 TESTOSTERONE CYPIONATE, TESTOSTERONE CYPIONATE
 TESTOSTERONE, TESTOSTERONE
 TRI-ESTARYLLA, ETHINYL ESTRADIOL
 TRI-LO-ESTARYLLA, ETHINYL ESTRADIOL
 VIENVA, ETHINYL ESTRADIOL
 VOLNEA, DESOGESTREL

XTTRIUM

* XTTRIUM LABORATORIES INC
 CHLORHEXIDINE GLUCONATE, CHLORHEXIDINE GLUCONATE
 EXIDINE, CHLORHEXIDINE GLUCONATE (OTC)

XTTRIUM LABS INC

* XTTRIUM LABORATORIES INC
 AMANTADINE HYDROCHLORIDE, AMANTADINE HYDROCHLORIDE
 LACTULOSE, LACTULOSE

**** Y ******YABAO PHARM**

* YABAO PHARMACEUTICAL CO LTD BEIJING
 GALANTAMINE HYDROBROMIDE, GALANTAMINE HYDROBROMIDE
 SORAFENIB TOSYLATE, SORAFENIB TOSYLATE

YAOPHARMA CO LTD

* YAOPHARMA CO LTD
 VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE

YICHANG HUMANWELL

* YICHANG HUMANWELL PHARMACEUTICAL CO LTD
 BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
 DESVENLAFAXINE SUCCINATE, DESVENLAFAXINE SUCCINATE
 FENOFIBRATE, FENOFIBRATE
 FENOFIBRIC ACID, CHOLINE FENOFIBRATE
 FLECAINIDE ACETATE, FLECAINIDE ACETATE
 GUANFACINE HYDROCHLORIDE, GUANFACINE HYDROCHLORIDE
 IBUPROFEN, IBUPROFEN
 MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 METOPROLOL SUCCINATE, METOPROLOL SUCCINATE
 NAPROXEN SODIUM, NAPROXEN SODIUM (OTC)
 POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
 VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE

YILING

* YILING PHARMACEUTICAL LTD
 ACYCLOVIR, ACYCLOVIR
 CELECOXIB, CELECOXIB
 CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE
 FELODIPINE, FELODIPINE
 LAMOTRIGINE, LAMOTRIGINE
 PAROXETINE HYDROCHLORIDE, PAROXETINE HYDROCHLORIDE
 PREGABALIN, PREGABALIN
 SIMVASTATIN, SIMVASTATIN
 VALACYCLOVIR HYDROCHLORIDE, VALACYCLOVIR HYDROCHLORIDE

YOUNGTECH PHARMS INC

* YOUNGTECH PHARMACEUTICALS INC
 METOPROLOL TARTRATE, METOPROLOL TARTRATE

YUNG SHIN PHARM

* YUNG SHIN PHARMACEUTICAL INDUSTRIAL CO LTD
 AZITHROMYCIN, AZITHROMYCIN
 CEFACLOR, CEFACLOR
 CEPHALEXIN, CEPHALEXIN

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** Y ****

- * YUNG SHIN PHARMACEUTICAL INDUSTRIAL CO LTD
CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
DICLOFENAC SODIUM AND MISOPROSTOL, DICLOFENAC SODIUM
ENTECAVIR, ENTECAVIR
FELODIPINE, FELODIPINE
MELOXICAM, MELOXICAM

**** Z ******ZEALAND PHARMA**

- * ZEALAND PHARMA US INC
ZEGALOGUE (AUTOINJECTOR), DASIGLUCAGON HYDROCHLORIDE
ZEGALOGUE, DASIGLUCAGON HYDROCHLORIDE

ZENNOVA

- * ZENNOVA PHARMACEUTICALS CHENGDU CO LTD
IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE
PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE

ZEVRA DENMARK

- * ZEVRA DENMARK AS
MIPLYFFA, ARIMOCLOMOL CITRATE

ZHEJIANG HISUN PHARM

- * ZHEJIANG HISUN PHARMACEUTICAL CO LTD
MYCOPHENOLATE MOFETIL, MYCOPHENOLATE MOFETIL

ZHEJIANG JINGXIN

- * ZHEJIANG JINGXIN PHARMACEUTICAL CO LTD
COLESEVELAM HYDROCHLORIDE, COLESEVELAM HYDROCHLORIDE
LEVETIRACETAM, LEVETIRACETAM

ZHEJIANG JIUZHOU

- * ZHEJIANG JIUZHOU BIOPHARMA CO LTD
CARBAMAZEPINE, CARBAMAZEPINE

ZHEJIANG JUTAI PHARM

- * ZHEJIANG JUTAI PHARMACEUTICAL CO LTD
AMANTADINE HYDROCHLORIDE, AMANTADINE HYDROCHLORIDE
BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
METOPROLOL SUCCINATE, METOPROLOL SUCCINATE

ZHEJIANG NOVUS PHARM

- * ZHEJIANG NOVUS PHARMACEUTICALS CO LTD
VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE

ZHEJIANG POLY PHARM

- * ZHEJIANG POLY PHARM CO LTD
AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE
FLUCONAZOLE, FLUCONAZOLE

ZHEJIANG XIANJU

- * ZHEJIANG XIANJU PHARMACEUTICAL CO LTD
PREDNISOLONE, PREDNISOLONE

ZHEJIANG YONGTAI

- * ZHEJIANG YONGTAI PHARMACEUTICAL CO LTD
ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM
GABAPENTIN, GABAPENTIN
ROSUVASTATIN CALCIUM, ROSUVASTATIN CALCIUM

ZHUHAI

- * ZHUHAI BEIHAI BIOTECH CO LTD
BEIZRAY, DOCETAXEL

ZMI PHARMA

- * ZMI PHARMA INC
ZIMHI, NALOXONE HYDROCHLORIDE

ZUREX PHARMA

- * ZUREX PHARMA
ZURAGARD, ISOPROPYL ALCOHOL (OTC)

ZYDUS

- * ZYDUS WORLDWIDE DMCC
AMANTADINE HYDROCHLORIDE, AMANTADINE HYDROCHLORIDE
MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE
OMEPRAZOLE AND SODIUM BICARBONATE, OMEPRAZOLE (OTC)
TOPIRAMATE, TOPIRAMATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** Z ****

* ZYDUS WORLDWIDE DMCC

VARENICLINE TARTRATE, VARENICLINE TARTRATE

ZYDUS HLTHCARE

* ZYDUS HEALTHCARE USA LLC

DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE

ISOSORBIDE MONONITRATE, ISOSORBIDE MONONITRATE

LANSOPRAZOLE, LANSOPRAZOLE

METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE

ZYDUS LIFESCIENCES

* ZYDUS LIFESCIENCES GLOBAL FZE

ACETAZOLAMIDE, ACETAZOLAMIDE

AZITHROMYCIN, AZITHROMYCIN

BACLOFEN, BACLOFEN

BALSALAZIDE DISODIUM, BALSALAZIDE DISODIUM

CHLORPROMAZINE HYDROCHLORIDE, CHLORPROMAZINE HYDROCHLORIDE

DAPSONE, DAPSONE

DOXORUBICIN HYDROCHLORIDE (LIPOSOMAL), DOXORUBICIN HYDROCHLORIDE

ERYTHROMYCIN, ERYTHROMYCIN

FLUDROCORTISONE ACETATE, FLUDROCORTISONE ACETATE

FLUPHENAZINE HYDROCHLORIDE, FLUPHENAZINE HYDROCHLORIDE

ICOSAPENT ETHYL, ICOSAPENT ETHYL

INDOMETHACIN, INDOMETHACIN

ISOSORBIDE DINITRATE, ISOSORBIDE DINITRATE

IVERMECTIN, IVERMECTIN

KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE

LEFLUNOMIDE, LEFLUNOMIDE

LIOTHYRONINE SODIUM, LIOTHYRONINE SODIUM

MECLIZINE HYDROCHLORIDE, MECLIZINE HYDROCHLORIDE

METHYLENE BLUE, METHYLENE BLUE

MIDODRINE HYDROCHLORIDE, MIDODRINE HYDROCHLORIDE

PHYTONADIONE, PHYTONADIONE

PROCHLORPERAZINE MALEATE, PROCHLORPERAZINE MALEATE

PROPAFENONE HYDROCHLORIDE, PROPAFENONE HYDROCHLORIDE

SILDENAFIL CITRATE, SILDENAFIL CITRATE

URSODIOL, URSODIOL

VALBENAZINE TOSYLATE, VALBENAZINE TOSYLATE

VIGABATRIN, VIGABATRIN

ZITUVIMET XR, METFORMIN HYDROCHLORIDE

ZITUVIMET, METFORMIN HYDROCHLORIDE

ZITUVIO, SITAGLIPTIN

* ZYDUS LIFESCIENCES LTD

ACYCLOVIR, ACYCLOVIR

BENAZEPRIL HYDROCHLORIDE, BENAZEPRIL HYDROCHLORIDE

BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE

BISOPROLOL FUMARATE, BISOPROLOL FUMARATE

CANDESARTAN CILEXETIL AND HYDROCHLOROTHIAZIDE, CANDESARTAN CILEXETIL

CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE

CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE

CYCLOPHOSPHAMIDE, CYCLOPHOSPHAMIDE

DARUNAVIR, DARUNAVIR

DESOXIMETASONE, DESOXIMETASONE

DEXAMETHASONE, DEXAMETHASONE

DICLOFENAC SODIUM, DICLOFENAC SODIUM

DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE

DOXEPIN HYDROCHLORIDE, DOXEPIN HYDROCHLORIDE

DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE

DUTASTERIDE, DUTASTERIDE

ESCITALOPRAM OXALATE, ESCITALOPRAM OXALATE

FEBUXOSTAT, FEBUXOSTAT

FELBAMATE, FELBAMATE

FLUOCINONIDE, FLUOCINONIDE

INDOMETHACIN, INDOMETHACIN

LEVOFLOXACIN, LEVOFLOXACIN

LIDOCAINE AND PRILOCAINE, LIDOCAINE

LOPERAMIDE HYDROCHLORIDE, LOPERAMIDE HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** Z ******* ZYDUS LIFESCIENCES LTD**

MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 METRONIDAZOLE, METRONIDAZOLE
 NYSTATIN AND TRIAMCINOLONE ACETONIDE, NYSTATIN
 NYSTATIN, NYSTATIN
 PIROXICAM, PIROXICAM
 ROPINIROLE HYDROCHLORIDE, ROPINIROLE HYDROCHLORIDE
 SUCRALFATE, SUCRALFATE
 TAVABOROLE, TAVABOROLE
 THEOPHYLLINE, THEOPHYLLINE
 TRETINOIN, TRETINOIN
 VALACYCLOVIR HYDROCHLORIDE, VALACYCLOVIR HYDROCHLORIDE
 VALSARTAN, VALSARTAN
 ZIPRASIDONE HYDROCHLORIDE, ZIPRASIDONE HYDROCHLORIDE
 ZONISAMIDE, ZONISAMIDE

ZYDUS PHARMS*** ZYDUS PHARMACEUTICALS USA INC**

ACAMPROSATE CALCIUM, ACAMPROSATE CALCIUM
 ACETAZOLAMIDE SODIUM, ACETAZOLAMIDE SODIUM
 ACETYLCYSTEINE, ACETYLCYSTEINE
 ACYCLOVIR SODIUM, ACYCLOVIR SODIUM
 ACYCLOVIR, ACYCLOVIR
 ADAPALENE AND BENZOYL PEROXIDE, ADAPALENE
 ALBENDAZOLE, ALBENDAZOLE
 ALBUTEROL SULFATE, ALBUTEROL SULFATE
 ALLOPURINOL, ALLOPURINOL
 AMANTADINE HYDROCHLORIDE, AMANTADINE HYDROCHLORIDE
 AMBRISENTAN, AMBRISENTAN
 AMITRIPTYLINE HYDROCHLORIDE, AMITRIPTYLINE HYDROCHLORIDE
 AMLODIPINE AND OLMESARTAN MEDOXOMIL, AMLODIPINE BESYLATE
 AMLODIPINE BESYLATE AND ATORVASTATIN CALCIUM, AMLODIPINE BESYLATE
 ARSENIC TRIOXIDE, ARSENIC TRIOXIDE
 ASPIRIN AND DIPYRIDAMOLE, ASPIRIN
 ATENOLOL AND CHLORTHALIDONE, ATENOLOL
 ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM
 AZELASTINE HYDROCHLORIDE, AZELASTINE HYDROCHLORIDE
 BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
 BORTEZOMIB, BORTEZOMIB
 BOSENTAN, BOSENTAN
 BUDESONIDE, BUDESONIDE
 BUMETANIDE, BUMETANIDE
 BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
 BUSPIRONE HYDROCHLORIDE, BUSPIRONE HYDROCHLORIDE
 CARBAMAZEPINE, CARBAMAZEPINE
 CARBIDOPA, CARBIDOPA
 CHLORPROMAZINE HYDROCHLORIDE, CHLORPROMAZINE HYDROCHLORIDE
 CHLORTHALIDONE, CHLORTHALIDONE
 CHOLESTYRAMINE LIGHT, CHOLESTYRAMINE
 CHOLESTYRAMINE, CHOLESTYRAMINE
 CISATRACURIUM BESYLATE, CISATRACURIUM BESYLATE
 CLINDAMYCIN PHOSPHATE AND BENZOYL PEROXIDE, BENZOYL PEROXIDE
 CLOBAZAM, CLOBAZAM
 CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
 CLOMIPRAMINE HYDROCHLORIDE, CLOMIPRAMINE HYDROCHLORIDE
 CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
 COLCHICINE, COLCHICINE
 COLESEVELAM HYDROCHLORIDE, COLESEVELAM HYDROCHLORIDE
 COLESTIPOL HYDROCHLORIDE, COLESTIPOL HYDROCHLORIDE
 CYANOCOBALAMIN, CYANOCOBALAMIN
 CYPROHEPTADINE HYDROCHLORIDE, CYPROHEPTADINE HYDROCHLORIDE
 DECITABINE, DECITABINE
 DEFERASIROX, DEFERASIROX
 DESMOPRESSIN ACETATE (NEEDS NO REFRIGERATION), DESMOPRESSIN ACETATE
 DESVENLAFAXINE SUCCINATE, DESVENLAFAXINE SUCCINATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** Z ****

* ZYDUS PHARMACEUTICALS USA INC
 DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE
 DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE
 DIFLUNISAL, DIFLUNISAL
 DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
 DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
 DOXAZOSIN MESYLATE, DOXAZOSIN MESYLATE
 DOXYCYCLINE, DOXYCYCLINE
 DOXYCYCLINE, DOXYCYCLINE HYCLATE
 DROXIDOPA, DROXIDOPA
 DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE
 DUTASTERIDE AND TAMSULOSIN HYDROCHLORIDE, DUTASTERIDE
 ELETRIPTAN HYDROBROMIDE, ELETRIPTAN HYDROBROMIDE
 EMTRICITABINE AND TENOFOVIR DISOPROXIL FUMARATE, EMTRICITABINE
 ENOXAPARIN SODIUM (PRESERVATIVE FREE), ENOXAPARIN SODIUM
 ENTECAVIR, ENTECAVIR
 ENZALUTAMIDE, ENZALUTAMIDE
 ERLOTINIB HYDROCHLORIDE, ERLOTINIB HYDROCHLORIDE
 ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM
 ESTRADIOL, ESTRADIOL
 ETHACRYNATE SODIUM, ETHACRYNATE SODIUM
 ETHINYL ESTRADIOL AND NORELGESTROMIN, ETHINYL ESTRADIOL
 ETODOLAC, ETODOLAC
 ETOMIDATE, ETOMIDATE
 EXEMESTANE, EXEMESTANE
 EZETIMIBE, EZETIMIBE
 FAMOTIDINE, FAMOTIDINE
 FESOTERODINE FUMARATE, FESOTERODINE FUMARATE
 FINGOLIMOD HYDROCHLORIDE, FINGOLIMOD HYDROCHLORIDE
 FLUCONAZOLE, FLUCONAZOLE
 FLUOCINONIDE, FLUOCINONIDE
 FULVESTRANT, FULVESTRANT
 GABAPENTIN, GABAPENTIN
 GLIPIZIDE, GLIPIZIDE
 GLYBURIDE AND METFORMIN HYDROCHLORIDE, GLYBURIDE
 GLYBURIDE, GLYBURIDE
 HALOPERIDOL DECANOATE, HALOPERIDOL DECANOATE
 IMATINIB MESYLATE, IMATINIB MESYLATE
 INDOMETHACIN, INDOMETHACIN
 ISOTRETINOIN, ISOTRETINOIN
 ITRACONAZOLE, ITRACONAZOLE
 IVABRADINE HYDROCHLORIDE, IVABRADINE HYDROCHLORIDE
 LABETALOL HYDROCHLORIDE, LABETALOL HYDROCHLORIDE
 LACOSAMIDE, LACOSAMIDE
 LAMOTRIGINE, LAMOTRIGINE
 LANSOPRAZOLE, LANSOPRAZOLE
 LENALIDOMIDE, LENALIDOMIDE
 LEVOTHYROXINE SODIUM, LEVOTHYROXINE SODIUM
 LINEZOLID, LINEZOLID
 LUBIPROSTONE, LUBIPROSTONE
 LURASIDONE HYDROCHLORIDE, LURASIDONE HYDROCHLORIDE
 MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
 MESALAMINE, MESALAMINE
 METHOTREXATE SODIUM, METHOTREXATE SODIUM
 METHYLPREDNISOLONE, METHYLPREDNISOLONE
 METOPROLOL SUCCINATE, METOPROLOL SUCCINATE
 MICAFUNGIN SODIUM, MICAFUNGIN SODIUM
 MIRABEGRON, MIRABEGRON
 MYCOPHENOLATE MOFETIL HYDROCHLORIDE, MYCOPHENOLATE MOFETIL HYDROCHLORIDE
 NADOLOL, NADOLOL
 NATEGLINIDE, NATEGLINIDE
 NELARABINE, NELARABINE
 NIFEDIPINE, NIFEDIPINE
 NITROFURANTOIN, NITROFURANTOIN, MACROCRYSTALLINE
 NITROGLYCERIN, NITROGLYCERIN

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** Z ****

* ZYDUS PHARMACEUTICALS USA INC
 NYSTATIN, NYSTATIN
 OLANZAPINE, OLANZAPINE
 OLMESARTAN MEDOXOMIL, OLMESARTAN MEDOXOMIL
 OMEPRAZOLE AND SODIUM BICARBONATE, OMEPRAZOLE
 OSELTAMIVIR PHOSPHATE, OSELTAMIVIR PHOSPHATE
 OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE
 PALIPERIDONE, PALIPERIDONE
 PEMETREXED DISODIUM, PEMETREXED DISODIUM
 PERPHENAZINE, PERPHENAZINE
 PHENTERMINE HYDROCHLORIDE, PHENTERMINE HYDROCHLORIDE
 PITAVASTATIN CALCIUM, PITAVASTATIN CALCIUM
 POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
 POTASSIUM CITRATE, POTASSIUM CITRATE
 PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE
 PYRIDOSTIGMINE BROMIDE, PYRIDOSTIGMINE BROMIDE
 RAMELTEON, RAMELTEON
 RIVASTIGMINE, RIVASTIGMINE
 ROFLUMILAST, ROFLUMILAST
 SACUBITRIL AND VALSARTAN, SACUBITRIL
 SCOPOLAMINE, SCOPOLAMINE
 SEVELAMER CARBONATE, SEVELAMER CARBONATE
 SIROLIMUS, SIROLIMUS
 SPIRONOLACTONE, SPIRONOLACTONE
 SUCCINYLCHOLINE CHLORIDE, SUCCINYLCHOLINE CHLORIDE
 TADALAFIL, TADALAFIL
 TAMOXIFEN CITRATE, TAMOXIFEN CITRATE
 TELMISARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 TELMISARTAN, TELMISARTAN
 TEMOZOLOMIDE, TEMOZOLOMIDE
 TERIFLUNOMIDE, TERIFLUNOMIDE
 TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE
 TRAZODONE HYDROCHLORIDE, TRAZODONE HYDROCHLORIDE
 TRIAMTERENE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 TRIAZOLAM, TRIAZOLAM
 TRIENTINE HYDROCHLORIDE, TRIENTINE HYDROCHLORIDE
 VARDENAFIL HYDROCHLORIDE, VARDENAFIL HYDROCHLORIDE
 VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE
 VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE
 VORICONAZOLE, VORICONAZOLE
 ZINC SULFATE, ZINC SULFATE
 ZOLMITRIPTAN, ZOLMITRIPTAN

ZYDUS PHARMS USA

* ZYDUS PHARMACEUTICALS USA INC
 AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
 ATENOLOL, ATENOLOL
 AZATHIOPRINE, AZATHIOPRINE
 BENZONATATE, BENZONATATE
 CLINDAMYCIN HYDROCHLORIDE, CLINDAMYCIN HYDROCHLORIDE
 HALOPERIDOL, HALOPERIDOL
 LAMOTRIGINE, LAMOTRIGINE
 MELOXICAM, MELOXICAM
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 NAPROXEN, NAPROXEN
 PAROXETINE HYDROCHLORIDE, PAROXETINE HYDROCHLORIDE
 PRAVASTATIN SODIUM, PRAVASTATIN SODIUM
 PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
 RAMIPRIL, RAMIPRIL
 RIBAVIRIN, RIBAVIRIN
 RISPERIDONE, RISPERIDONE
 SIMVASTATIN, SIMVASTATIN
 VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE
 WARFARIN SODIUM, WARFARIN SODIUM

ZYDUS PHARMS USA INC

* ZYDUS PHARMACEUTICALS USA INC

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** Z ******* ZYDUS PHARMACEUTICALS USA INC**

AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE
 ANASTROZOLE, ANASTROZOLE
 ATOMOXETINE HYDROCHLORIDE, ATOMOXETINE HYDROCHLORIDE
 BROMOCRIPTINE MESYLATE, BROMOCRIPTINE MESYLATE
 CARVEDILOL, CARVEDILOL
 DIPYRIDAMOLE, DIPYRIDAMOLE
 DIVALPROEX SODIUM, DIVALPROEX SODIUM
 FINASTERIDE, FINASTERIDE
 GABAPENTIN, GABAPENTIN
 GALANTAMINE HYDROBROMIDE, GALANTAMINE HYDROBROMIDE
 GLIPIZIDE AND METFORMIN HYDROCHLORIDE, GLIPIZIDE
 HYDROXYCHLOROQUINE SULFATE, HYDROXYCHLOROQUINE SULFATE
 IRBESARTAN, IRBESARTAN
 LAMOTRIGINE, LAMOTRIGINE
 LEVETIRACETAM, LEVETIRACETAM
 LOSARTAN POTASSIUM AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 LOSARTAN POTASSIUM, LOSARTAN POTASSIUM
 OMEPRAZOLE, OMEPRAZOLE
 PIOGLITAZONE HYDROCHLORIDE, PIOGLITAZONE HYDROCHLORIDE
 PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE
 PROPRANOLOL HYDROCHLORIDE, PROPRANOLOL HYDROCHLORIDE
 RISPERIDONE, RISPERIDONE
 TAMSULOSIN HYDROCHLORIDE, TAMSULOSIN HYDROCHLORIDE
 TOPIRAMATE, TOPIRAMATE
 TRAMADOL HYDROCHLORIDE AND ACETAMINOPHEN, ACETAMINOPHEN
 TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE
 VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE
 ZOLMITRIPTAN, ZOLMITRIPTAN

ZYLA*** ZYLA LIFE SCIENCES US INC**

INDOCIN, INDOMETHACIN
 SPRIX, KETOROLAC TROMETHAMINE

45TH EDITION - 2025 - APPROVED DRUG PRODUCT LIST

APPENDIX C**UNIFORM TERMS*****DOSAGE FORMS***

AEROSOL, FOAM	POWDER
AEROSOL, METERED	POWDER, EXTENDED RELEASE
CAPSULE	POWDER, METERED
CAPSULE, DELAYED REL PELLETS	RING
CAPSULE, DELAYED RELEASE	SHAMPOO
CAPSULE, EXTENDED RELEASE	SOLUTION
CAPSULE, PELLETS	SOLUTION FOR SLUSH
CAPSULE, TABLET	SOLUTION, EXTENDED RELEASE
CAPSULE, TABLET, TABLET	SOLUTION, GEL FORMING/DROPS
CLOTH	SOLUTION, METERED
CONCENTRATE	SOLUTION/DROPS
CREAM	SPONGE
CREAM, AUGMENTED	SPRAY
CREAM, INSERT	SPRAY, METERED
ELIXIR	SUPPOSITORY
EMULSION	SUSPENSION
ENEMA	SUSPENSION, EXTENDED RELEASE
FILM	SUSPENSION, LIPOSOMAL
FILM, EXTENDED RELEASE	SUSPENSION/DROPS
FOAM	SWAB
FOR SOLUTION	SYRUP
FOR SUSPENSION	SYSTEM
FOR SUSPENSION, DELAYED RELEASE	TABLET
FOR SUSPENSION, EXTENDED RELEASE	TABLET, CHEWABLE
GAS	TABLET, DELAYED RELEASE
GEL	TABLET, EFFERVESCENT
GEL, AUGMENTED	TABLET, EXTENDED RELEASE
GEL, METERED	TABLET, EXTENDED RELEASE,
GRANULE	CHEWABLE
GRANULE, DELAYED RELEASE	TABLET, FOR SUSPENSION
GRANULES	TABLET, ORALLY DISINTEGRATING
GUM, CHEWING	TABLET, ORALLY DISINTEGRATING,
IMPLANT	DELAYED RELEASE
INJECTABLE	TABLET, ORALLY DISINTEGRATING,
INJECTABLE, LIPID COMPLEX	EXTENDED RELEASE
INJECTABLE, LIPOSOMAL	TAPE
INSERT	TROCHE/LOZENGE
INSERT, EXTENDED RELEASE	
JELLY	
LIQUID	
LOTION	
LOTION, AUGMENTED	
LOTION/SHAMPOO	
OIL	
OIL/DROPS	
OINTMENT	
OINTMENT, AUGMENTED	
PASTE	
PATCH	
PELLET	
PELLETS	

Note: Terms comprise currently marketed products

APPENDIX C**UNIFORM TERMS*****ROUTES OF ADMINISTRATION***

BUCCAL	OPHTHALMIC
CARDIAC	ORAL
DENTAL	ORAL-21
ENDOCERVICAL	ORAL-28
ENDOTRACHEAL	OTIC
ENTERAL	PERFUSION
IMPLANTATION	PERIARTICULAR
INHALATION	PERIODONTAL
INJECTION	PYELOCALYCEAL
INTERSTITIAL	RECTAL
INTRA-ANAL	SPINAL
INTRA-ARTERIAL	SUBCUTANEOUS
INTRA-ARTICULAR	SUBLINGUAL
INTRACAMERAL	TOPICAL
INTRACAVITARY	TRANSDERMAL
INTRACRANIAL	URETHRAL
INTRADERMAL	VAGINAL
INTRAMUSCULAR	
INTRAOCULAR	
INTRAOSSEOUS	
INTRAPERITONEAL	
INTRAPLEURAL	
INTRATHECAL	
INTRAUTERINE	
INTRAVENOUS	
INTRAVESICAL	
INTRAVITREAL	
IRRIGATION	
N/A	
NASAL	

APPENDIX C**UNIFORM TERMS*****ABBREVIATIONS***

AMP	AMPULE
AMPICIL	AMPICILLIN
APPROX	APPROXIMATELY
BOT	BOTTLE
CI	CURIE
CSR	CAROTID SINUS REFLEX
CU	CLINICAL UNITS
DIPROP	DIPROPIONATE
ELECT	ELECTROLYTE
EQ	EQUIVALENT TO
ER	EXTENDED RELEASE
GM	GRAM
HBR	HYDROBROMIDE
HCL	HYDROCHLORIDE
HR	HOUR
IM	INTRAMUSCULAR
INH	INHALATION
IU	INTERNATIONAL UNITS
IV	INTRAVENOUS
KIU	KALLIKREIN INHIBITOR UNITS
MCG	MICROGRAM
MCI	MILLICURIE
MEQ	MILLIEQUIVALENT
MG	MILLIGRAM
ML	MILLILITER
N/A	NOT APPLICABLE
PPM	PARTS PER MILLION
REL	RELEASE
SC	SUBCUTANEOUS
SQ CM	SQUARE CENTIMETER
U	UNITS
UCI	MICROCURIE
UMOLAR	MICROMOLAR
USP	UNITED STATES PHARMACOPEIA

PATENT AND EXCLUSIVITY INFORMATION ADDENDUM

This *Addendum* identifies drugs that qualify under the Federal Food, Drug, and Cosmetic Act (the FD&C Act) for periods of exclusivity and provides patent information that has been submitted to the Food and Drug Administration (FDA) concerning the listed drug products.

Exclusivity

This *Addendum* identifies:

- Drugs approved under Section 505(c) of the FD&C Act that have qualified under the Drug Price Competition and Patent Term Restoration Act of 1984 (Hatch-Waxman Amendments) for 5-year and 3-year periods of exclusivity pursuant to Section 505(c)(3)(E) and Section 505(j)(5)(F) of the FD&C Act
- Drugs that have qualified for Orphan Drug Exclusivity pursuant to Section 527 of the FD&C Act
- Drugs that have qualified for Pediatric Exclusivity pursuant to Section 505A of the FD&C Act
- Drugs that have qualified for Generating Antibiotic Incentives Now (GAIN) exclusivity pursuant to Section 505E of the FD&C Act
- Generic drugs approved under Section 505(j) of the FD&C Act that have qualified for 180-day exclusivity pursuant to Section 505(j)(5)(B)(iv) of the FD&C Act
- Generic drugs approved under Section 505(j) of the FD&C Act that have qualified for Competitive Generic Therapy (CGT) exclusivity pursuant to Section 505(j)(5)(B)(v) of the FD&C Act

This section is arranged in alphabetical order by name of the active ingredient, followed by the proprietary name (brand name or trade name) of the drug product. Active ingredient headings for multiple active ingredient, fixed-combination drug products are arranged alphabetically.

Individual descriptions of the protected use have been added to each Orphan Drug Exclusivity entry listed in the Orange Book. Such descriptions of Orphan Drug Exclusivity were included beginning with the 38th edition of the Orange Book. The ODE* code means that the timing of approval of certain follow-on applications may be subject to delay due to ODE for another drug that has the same active moiety.

For an explanation of the codes used in the *Addendum*, see the *Patent and Exclusivity Terms* section. The exclusivity codes are general shorthand descriptions and do not necessarily identify, with specificity, the actual scope of exclusivity.

Patent Information

The FD&C Act requires that certain patent information be filed with all newly submitted new drug applications (NDA) and with certain supplements to approved NDAs. Form FDA 3542a must be used for this pre-approval submission.¹

The FD&C Act also requires that certain patent information be filed after approval of an NDA. To be considered timely filed, such patent information must be submitted according to the following timeframes:

- not later than 30 days after the date of approval of the NDA² or supplement; or
- not later than 30 days after the date of issuance of the patent when a patent is issued after approval of the NDA or supplement.

Form FDA 3542 must be used for these submissions.³ FDA publishes certain information from Form FDA 3542 in the Orange Book.

The Orange Book includes the patent submission date (i.e., the date on which FDA receives patent information from the NDA holder on Form FDA 3542) for each newly listed patent to facilitate assessments of whether patent information is untimely filed with respect to a pending 505(b)(2) application or abbreviated new drug application (ANDA) and whether patent information was submitted before the date on which a 505(b)(2) application or ANDA (excluding an amendment or supplement to the 505(b)(2) application or ANDA) was submitted.⁴

Section 505(c)(2) of the FD&C Act requires, with reference to Section 505(b)(1)(A)(viii) of the FD&C Act, that applicants submit certain patent information for each patent for which a claim of patent infringement could reasonably be asserted if a person not licensed by the owner of the patent engaged in the manufacture, use, or sale of the drug and that:

- claims the drug for which the applicant submitted the application and is a drug substance (active ingredient) patent or a drug product (formulation or composition) patent; or
- claims a method of using such drug for which approval has been granted in the application.

This information, as provided by the NDA holder on Form FDA 3542, will be published in the Orange Book, as described above. An NDA holder submitting information on a patent that claims both the drug substance and the drug product (and is eligible for listing on either basis) is required only to specify that the patent claims either the drug substance or the drug product.⁵

The Addendum lists patent and exclusivity information up to January of the Edition year. The monthly Cumulative Supplements to the Annual Edition list patent and exclusivity information changes since the Annual Edition Addendum. Since all parts of this publication are subject to changes, additions, or

¹ 21 CFR 314.53(c)(2)(i); 21 CFR 314.53(d).

² Please note that the date of approval for an NDA for a drug for which FDA intends to recommend controls under the Controlled Substances Act is the later of the date on the approval letter for the NDA or the date of issuance of the interim final rule controlling the drug (see Section 505(x)(1) and (2) of the FD&C Act).

³ 21 CFR 314.53(c)(2)(ii); 21 CFR 314.53(d).

⁴ See 21 CFR 314.50(i)(4) and 314.94(a)(12)(vi); see also 21 CFR 314.107(b)(3)(i). The submission date for patent information is determined in accordance with 21 CFR 314.53(d)(5).

⁵ 21 CFR 314.53.

deletions, the [Orange Book](#), updated regularly, should be consulted for the most recent patent and exclusivity information.

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>ABACAVIR SULFATE; DOLUTEGRAVIR SODIUM; LAMIVUDINE - TRIUMEO</u>						
N 205551	001	8129385	Oct 05, 2027	DS DP	M-294	Jun 15, 2026
		8129385*PED	Apr 05, 2028		PED	Dec 15, 2026
		9242986	Dec 08, 2029	DS DP		
		9242986*PED	Jun 08, 2030			
<u>ABACAVIR SULFATE; DOLUTEGRAVIR SODIUM; LAMIVUDINE - TRIUMEO PD</u>						
N 215413	001	8129385	Oct 05, 2027	DS DP	NPP	Jun 15, 2026
		8129385*PED	Apr 05, 2028		PED	Dec 15, 2026
		9242986	Dec 08, 2029	DS DP		
		9242986*PED	Jun 08, 2030			
<u>ABALOPARATIDE - TYMLOS</u>						
N 208743	001	10996208	Apr 30, 2038	DP	I-907	Dec 19, 2025
		11255842	Jan 10, 2040	U-3322		
		11680942	Jan 10, 2040	U-3322		
		11782041	Apr 30, 2038	DP U-2009		
		11782041	Apr 30, 2038	DP U-3543		
		11977067	Apr 30, 2038	DP		
		7803770	Apr 28, 2031	U-2009		
		8148333	Nov 08, 2027	DP		
		8748382	Oct 03, 2027	U-2009		
		8748382	Oct 03, 2027	U-3543		
		RE49444	Apr 28, 2031	U-2009		
		RE49444	Apr 28, 2031	U-3543		
<u>ABAMETAPIR - XEGLYZE</u>						
N 206966	001	10292389	Dec 17, 2034	DP U-2863	NCE	Jul 24, 2025
		7812163	Oct 28, 2026	DP U-2863		
<u>ABEMACICLIB - VERZENIO</u>						
N 208716	001	7855211	Sep 28, 2031	DS DP U-2135		
		7855211	Sep 28, 2031	DS DP U-3242		
		7855211	Sep 28, 2031	DS DP U-3265		
		7855211	Sep 28, 2031	DS DP U-3546		
<u>ABEMACICLIB - VERZENIO</u>						
N 208716	002	7855211	Sep 28, 2031	DS DP U-2135		
		7855211	Sep 28, 2031	DS DP U-3242		
		7855211	Sep 28, 2031	DS DP U-3265		
		7855211	Sep 28, 2031	DS DP U-3546		
<u>ABEMACICLIB - VERZENIO</u>						
N 208716	003	7855211	Sep 28, 2031	DS DP U-2135		
		7855211	Sep 28, 2031	DS DP U-3242		
		7855211	Sep 28, 2031	DS DP U-3265		
		7855211	Sep 28, 2031	DS DP U-3546		
<u>ABEMACICLIB - VERZENIO</u>						
N 208716	004	7855211	Sep 28, 2031	DS DP U-1981		
		7855211	Sep 28, 2031	DS DP U-2135		
		7855211	Sep 28, 2031	DS DP U-3242		
		7855211	Sep 28, 2031	DS DP U-3265		
		7855211	Sep 28, 2031	DS DP U-3546		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>ABIRATERONE ACETATE - YONSA</u>						
N 210308	001 10292990	May 20, 2034	U-2535			
	9889144	Mar 17, 2034	DP			
<u>ABIRATERONE ACETATE; NIRAPARIB TOSYLATE - AKEEGA</u>						
N 216793	001 11091459	Mar 27, 2038	DS DP		NP	Aug 11, 2026
	11207311	Jul 28, 2037	U-2830			
	11673877	Mar 27, 2038	DS DP			
	11986468	Jul 28, 2037	U-2830			
	11986469	Jul 28, 2037	U-2830			
	11992486	Jul 28, 2037	U-2830			
	8071579	Aug 12, 2027	U-2830			
	8071623	Mar 27, 2031	DS DP			
	8143241	Aug 12, 2027	U-2830			
	8436185	Apr 24, 2029	DS DP			
	8859562	Aug 04, 2031	U-2830			
<u>ABIRATERONE ACETATE; NIRAPARIB TOSYLATE - AKEEGA</u>						
N 216793	002 11091459	Mar 27, 2038	DS DP		NP	Aug 11, 2026
	11207311	Jul 28, 2037	U-2830			
	11673877	Mar 27, 2038	DS DP			
	11986468	Jul 28, 2037	U-2830			
	11986469	Jul 28, 2037	U-2830			
	11992486	Jul 28, 2037	U-2830			
	8071579	Aug 12, 2027	U-2830			
	8071623	Mar 27, 2031	DS DP			
	8143241	Aug 12, 2027	U-2830			
	8436185	Apr 24, 2029	DS DP			
	8859562	Aug 04, 2031	U-2830			
<u>ABROCITINIB - CIBINOO</u>						
N 213871	001 9035074	Feb 19, 2034	DS DP		NCE	Jan 14, 2027
	9545405	Feb 19, 2034	DS DP		NPP	Feb 09, 2026
	9549929	Feb 19, 2034	U-3195			
<u>ABROCITINIB - CIBINOO</u>						
N 213871	002 9035074	Feb 19, 2034	DS DP		NCE	Jan 14, 2027
	9545405	Feb 19, 2034	DS DP		NPP	Feb 09, 2026
	9549929	Feb 19, 2034	U-3195			
<u>ABROCITINIB - CIBINOO</u>						
N 213871	003 9035074	Feb 19, 2034	DS DP		NCE	Jan 14, 2027
	9545405	Feb 19, 2034	DS DP		NPP	Feb 09, 2026
	9549929	Feb 19, 2034	U-3195			
<u>ACALABRUTINIB - CALQUENCE</u>						
N 210259	001 10167291	Jul 01, 2036	DP U-2145		ODE-274	Nov 21, 2026
	10167291	Jul 01, 2036	DP U-2666			
	10167291	Jul 01, 2036	DP U-2667			
	10167291	Jul 01, 2036	DP U-2668			
	10167291	Jul 01, 2036	DP U-2669			
	10167291	Jul 01, 2036	DP U-2670			
	10167291	Jul 01, 2036	DP U-2671			
	10239883	Jul 11, 2032	U-2666			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>ACALABRUTINIB - CALOQUENCE</u>						
N 210259 001	10239883	Jul 11, 2032				U-2668
	10272083	Jan 21, 2035				U-2519
	10272083	Jan 21, 2035				U-2682
	10272083	Jan 21, 2035				U-2683
	10272083	Jan 21, 2035				U-2684
	10272083	Jan 21, 2035				U-2685
	10272083	Jan 21, 2035				U-2686
	10272083	Jan 21, 2035				U-2687
	11771696	Jan 21, 2035				U-3710
	7459554	Nov 24, 2026	DS			
	9290504	Jul 11, 2032	DS DP			
	9758524	Jul 11, 2032				U-2145
	9796721	Jul 01, 2036	DS DP			U-2145
	9796721	Jul 01, 2036	DS DP			U-2666
	9796721	Jul 01, 2036	DS DP			U-2667
	9796721	Jul 01, 2036	DS DP			U-2668
	9796721	Jul 01, 2036	DS DP			U-2669
	9796721	Jul 01, 2036	DS DP			U-2670
	9796721	Jul 01, 2036	DS DP			U-2671
<u>ACALABRUTINIB MALEATE - CALOQUENCE</u>						
N 216387 001	10239883	Jul 11, 2032				U-2666
	10239883	Jul 11, 2032				U-2668
	10272083	Jan 21, 2035				U-2519
	10272083	Jan 21, 2035				U-2682
	10272083	Jan 21, 2035				U-2683
	10272083	Jan 21, 2035				U-2684
	10272083	Jan 21, 2035				U-2685
	10272083	Jan 21, 2035				U-2686
	10272083	Jan 21, 2035				U-2687
	11059829	Jul 01, 2036	DS DP			U-2145
	11059829	Jul 01, 2036	DS DP			U-2666
	11059829	Jul 01, 2036	DS DP			U-2667
	11059829	Jul 01, 2036	DS DP			U-2668
	11059829	Jul 01, 2036	DS DP			U-2669
	11059829	Jul 01, 2036	DS DP			U-2670
	11059829	Jul 01, 2036	DS DP			U-2671
	11771696	Jan 21, 2035				U-3710
	7459554	Nov 24, 2026	DS			
	9290504	Jul 11, 2032	DS DP			
	9758524	Jul 11, 2032				U-2145
<u>ACETAMINOPHEN - OFIRMEV</u>						
N 022450 001	10383834	Nov 13, 2028				U-2262
	10383834	Nov 13, 2028				U-2621
	9399012	Sep 11, 2031				U-2261
	9399012	Sep 11, 2031				U-2262
	9399012*PED	Mar 11, 2032				
	9610265	Nov 13, 2028				U-2263
	9610265*PED	May 13, 2029				
	9987238	Nov 13, 2028				U-2261

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>ACETAMINOPHEN - OFIRMEV</u>						
N 022450	001 9987238*PED	May 13, 2029				
<u>ACETAMINOPHEN - ACETAMINOPHEN</u>						
N 204767	001 8741959	Apr 19, 2030	DP			
<u>ACETAMINOPHEN; BENZHYDROCODONE HYDROCHLORIDE - APADAZ</u>						
N 208653	001 8461137	Feb 22, 2031	DS DP			
	8748413	Jul 01, 2030	DS DP			
	8828978	Jul 01, 2030	DP			
	9132125	Jul 01, 2030	DS DP U-2249			
	9549923	Jul 01, 2030	DS DP			
<u>ACETAMINOPHEN; BENZHYDROCODONE HYDROCHLORIDE - APADAZ</u>						
N 208653	002 8461137	Feb 22, 2031	DS DP			
	8748413	Jul 10, 2030	DS DP			
	8828978	Jul 01, 2030	DP			
	9132125	Jul 01, 2030	DS DP U-2249			
	9549923	Jul 01, 2030	DS DP			
<u>ACETAMINOPHEN; BENZHYDROCODONE HYDROCHLORIDE - APADAZ</u>						
N 208653	003 8461137	Feb 22, 2031	DS DP			
	8748413	Jul 10, 2030	DS DP			
	8828978	Jul 01, 2030	DP			
	9132125	Jul 01, 2030	DS DP U-2249			
	9549923	Jul 01, 2030	DS DP			
<u>ACETAMINOPHEN; IBUPROFEN - COMBOGESIC</u>						
N 209471	001 10532036	Sep 22, 2025	U-3553		NP	Mar 01, 2026
	11197830	Feb 27, 2039	DP U-3553			
	11534407	Feb 27, 2039	DP U-3553			
<u>ACETAMINOPHEN; IBUPROFEN - ADVIL DUAL ACTION WITH ACETAMINOPHEN</u>						
N 211733	001 11918693	Jul 09, 2041	DP			
<u>ACETAMINOPHEN; IBUPROFEN SODIUM - COMBOGESIC IV</u>						
N 215320	001 11213498	Jan 14, 2036	DP		NP	Oct 17, 2026
	11389416	Jul 17, 2035	DP			
	11446266	Oct 26, 2031	U-3744			
	11446266	Oct 26, 2031	U-3745			
	11896567	Oct 26, 2031	U-3744			
	11896567	Oct 26, 2031	U-3745			
	12083087	Jul 17, 2035	U-3744			
	12083087	Jul 17, 2035	U-3745			
<u>ACETAMINOPHEN; OXYCODONE HYDROCHLORIDE - XARTEMIS XR</u>						
N 204031	001 7976870	Jun 01, 2027	U-1498			
	8372432	Mar 11, 2029	DP U-1499			
	8377453	Nov 19, 2029	DP U-1499			
	8394408	Mar 11, 2029	DP			
	8597681	Dec 21, 2030	DP			
	8658631	May 16, 2032	DP			
	8668929	Mar 11, 2029	U-1499			
	8741885	May 16, 2032	DP U-1499			
	8980319	Dec 21, 2030	DP			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>ACETAMINOPHEN; OXYCODONE HYDROCHLORIDE - XARTEMIS XR</u>						
N 204031	001 8992975	May 16, 2032	DP			
	9050335	May 16, 2032	DP			
	9468636	May 16, 2032			U-1499	
<u>ACETYLCYSTEINE - ACETADOTE</u>						
N 021539	001 8148356	May 21, 2026	DP			
	8399445	Aug 24, 2025			U-1373	
	8653061	Aug 24, 2025			U-1373	
	8722738	Apr 06, 2032			U-1373	
	9327028	Jul 21, 2031			U-1839	
<u>ACETYLCYSTEINE - CETYLEV</u>						
N 207916	001 8747894	May 08, 2032	DP		U-1373	
	9427421	May 08, 2032	DP			
	9561204	May 08, 2032			U-1373	
<u>ACETYLCYSTEINE - CETYLEV</u>						
N 207916	002 8747894	May 08, 2032	DP		U-1373	
	9427421	May 08, 2032	DP			
	9561204	May 08, 2032			U-1373	
<u>ACETYLCYSTEINE LYSINE - LEGUBETI</u>						
N 215040	001				NP	Feb 13, 2027
<u>ACETYLCYSTEINE LYSINE - LEGUBETI</u>						
N 215040	002				NP	Feb 13, 2027
<u>ACLIDINIUM BROMIDE - TUDORZA PRESSAIR</u>						
N 202450	001 10085974	Mar 13, 2029	DP		U-2513	
	11000517	Mar 13, 2029	DP		U-2513	
	8051851	Apr 22, 2027	DP			
	RE46417	Feb 10, 2025	DS DP		U-2513	
<u>ACLIDINIUM BROMIDE; FORMOTEROL FUMARATE - DUAKLIR PRESSAIR</u>						
N 210595	001 10085974	Mar 13, 2029	DP		U-2513	
	11000517	Mar 13, 2029	DP		U-2513	
	8051851	Apr 22, 2027	DP			
	RE46417	Feb 10, 2025	DS DP		U-2513	
<u>ACORAMIDIS HYDROCHLORIDE - ATTRUBY</u>						
N 216540	001 10398681	May 05, 2031	DP		NCE	Nov 22, 2029
	10513497	Feb 16, 2038	DS			
	10842777	May 05, 2031			U-4046	
	11058668	Mar 22, 2039			U-4046	
	11260047	Aug 16, 2039	DP			
	11919865	Feb 16, 2038			U-4046	
	12005043	Aug 16, 2039	DP		U-4046	
	12070449	Mar 22, 2039			U-4046	
	8877795	May 05, 2031	DP			
	9169214	May 05, 2031	DP		U-4046	
	9642838	May 05, 2031	DS DP			
	9913826	Mar 14, 2033			U-4046	

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>ACYCLOVIR - AVACLYR</u>						
N 202408	001				ODE-235	Mar 29, 2026
<u>ACYCLOVIR - SITAVIG</u>						
N 203791	001	8592434	Jun 16, 2030	DP U-1460		
		8747896	Jun 03, 2027	DP U-1460		
		8791127	Mar 23, 2027	DP U-1460		
<u>ADAGRASIB - KRAZATI</u>						
N 216340	001	10689377	May 17, 2037	DS DP U-3490	NCE	Dec 12, 2027
		10689377	May 17, 2037	DS DP U-3953	ODE-352	Dec 12, 2029
<u>ADAPALENE - DIFFERIN</u>						
N 021753	001	7579377	Feb 23, 2025	U-818		
<u>ADAPALENE - DIFFERIN</u>						
N 022502	001	7998467	May 31, 2028	DP U-1078		
		8435502	Sep 15, 2026	DP U-1078		
		8709392	Sep 15, 2026	DP U-1078		
<u>ADAPALENE; BENZOYL PEROXIDE - EPIDUO</u>						
N 022320	001	7820186	Nov 23, 2025	DP		
		8071644	Jul 18, 2027	DP U-1078		
		8080537	Jul 18, 2027	U-1078		
		8129362	Jul 18, 2027	U-1078		
<u>ADAPALENE; BENZOYL PEROXIDE; CLINDAMYCIN PHOSPHATE - CABTREO</u>						
N 216632	001	10220049	Jun 03, 2029	DP U-3713	NP	Oct 20, 2026
		10624918	Jun 03, 2029	U-3713		
		11389467	Dec 28, 2040	DP U-3713		
		12128059	Jul 31, 2040	DP U-3713		
		12133859	Jul 31, 2040	DP		
		12138278	Jul 31, 2040	DP		
		8288434	Aug 05, 2029	DP U-3713		
		9561208	Jun 03, 2029	DP U-3713		
<u>AFAMELANOTIDE - SCENESSE</u>						
N 210797	001	10076555	Feb 11, 2025	U-2638	ODE-270	Oct 08, 2026
		8334265	Mar 11, 2029	U-2638		
<u>AFATINIB DIMALEATE - GILOTRIF</u>						
N 201292	001	10004743	Jul 05, 2030	DP	M-276	Apr 07, 2025
		10004743*PED	Jan 05, 2031		ODE-230	Jan 12, 2025
		8426586	Oct 10, 2029	DS	PED	Jul 12, 2025
		8426586*PED	Apr 10, 2030		PED	Oct 07, 2025
		8545884	Dec 19, 2029	DP		
		8545884*PED	Jun 19, 2030			
		9539258	Nov 09, 2026	U-1950		
		9539258*PED	May 09, 2027			
		RE43431	Jan 13, 2026	DS		
		RE43431*PED	Jul 13, 2026			
<u>AFATINIB DIMALEATE - GILOTRIF</u>						
N 201292	002	10004743	Jul 05, 2030	DP	M-276	Apr 07, 2025
		10004743*PED	Jan 05, 2031		ODE-230	Jan 12, 2025
		8426586	Oct 10, 2029	DS	PED	Jul 12, 2025

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>AFATINIB DIMALEATE - GILOTRIF</u>						
N 201292 002	8426586*PED	Apr 10, 2030			PED	Oct 07, 2025
	8545884	Dec 19, 2029	DP			
	8545884*PED	Jun 19, 2030				
	9539258	Nov 09, 2026		U-1950		
	9539258*PED	May 09, 2027				
	RE43431	Jan 13, 2026	DS			
	RE43431*PED	Jul 13, 2026				
<u>AFATINIB DIMALEATE - GILOTRIF</u>						
N 201292 003	10004743	Jul 05, 2030	DP		M-276	Apr 07, 2025
	10004743*PED	Jan 05, 2031			ODE-230	Jan 12, 2025
	8426586	Oct 10, 2029	DS		PED	Jul 12, 2025
	8426586*PED	Apr 10, 2030			PED	Oct 07, 2025
	8545884	Dec 19, 2029	DP			
	8545884*PED	Jun 19, 2030				
	9539258	Nov 09, 2026		U-1950		
	9539258*PED	May 09, 2027				
	RE43431	Jan 13, 2026	DS			
	RE43431*PED	Jul 13, 2026				
<u>AIR POLYMER-TYPE A - EXEM FOAM KIT</u>						
N 212279 001	9034300	Oct 15, 2030	DP	U-2663		
	9259494	May 04, 2035	DP	U-2663		
	9849199	Feb 11, 2036	DP			
<u>ALBUTEROL SULFATE - PROAIR HEFA</u>						
N 021457 001	10022509	May 18, 2031	DP			
	10022510	May 18, 2031	DP			
	10086156	May 18, 2031	DP			
	10561808	Jan 01, 2032	DP			
	10695512	May 18, 2031	DP			
	11395889	May 18, 2031	DP			
	8132712	Sep 07, 2028	DP			
	9463289	May 18, 2031	DP			
	9808587	May 18, 2031	DP			
<u>ALBUTEROL SULFATE - PROAIR RESPICLICK</u>						
N 205636 001	10022510	May 18, 2031	DP			
	10124131	May 18, 2031	DP			
	10561808	Jan 01, 2032	DP			
	10765820	May 19, 2025	DP			
	8651103	Mar 26, 2028	DP			
	8978966	Jan 13, 2032	DP			
	9216260	Jun 28, 2031	DP			
	9463288	May 19, 2025	DP			
	9731087	May 18, 2031	DP			
<u>ALBUTEROL SULFATE - PROAIR DIGIHALER</u>						
N 205636 002	10022510	May 18, 2031	DP			
	10124131	May 18, 2031	DP			
	10561808	Jan 01, 2032	DP			
	10569034	Aug 16, 2036	DP			
	10765820	May 19, 2025	DP			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>ALBUTEROL SULFATE - PROAIR DIGIHALER</u>						
N 205636 002	10918816	Dec 14, 2035	DP			
	11000653	Dec 18, 2038	DP			
	11173259	Jul 06, 2040	DP			
	11266796	Feb 22, 2041	DP			
	11344685	Sep 26, 2039	DP			
	11351317	Feb 10, 2038	DP			
	11357935	Sep 24, 2038	DP			
	11439777	May 24, 2040	DP			
	11464923	Jun 19, 2040	DP			
	8651103	Mar 26, 2028	DP			
	8978966	Jan 13, 2032	DP			
	9216260	Jun 28, 2031	DP			
	9463288	May 19, 2025	DP			
	9731087	May 18, 2031	DP			
	9782550	Aug 28, 2035	DP			
	9782551	Aug 28, 2035	DP			
<u>ALBUTEROL SULFATE; BUDESONIDE - AIRSUPRA</u>						
N 214070 001	9415009	May 28, 2030	U-3509		NP	Jan 10, 2026
<u>ALBUTEROL SULFATE; IPRATROPIUM BROMIDE - COMBIVENT RESPIMAT</u>						
N 021747 001	7396341	Oct 10, 2026	DP			
	7837235	Mar 13, 2028	DP			
	7896264	May 26, 2025	DP			
	8733341	Oct 16, 2030	DP			
	9027967	Mar 31, 2027	DP			
<u>ALCAFTADINE - LASTACRAFT</u>						
N 022134 001	10617695	Mar 19, 2027	DP U-3267			
	8664215	Dec 23, 2027	U-3267			
<u>ALCOHOL - ABLYSINOL</u>						
N 207987 001					ODE-192	Jun 21, 2025
<u>ALCOHOL - ABLYSINOL</u>						
N 207987 002					ODE-192	Jun 21, 2025
<u>ALECTINIB HYDROCHLORIDE - ALECENSA</u>						
N 208434 001	10350214	Apr 24, 2035	DP		I-947	Apr 18, 2027
	11433076	Apr 24, 2035	DP		ODE-477	Apr 18, 2031
	9126931	May 29, 2031	DS			
	9365514	Mar 04, 2032	DP			
	9440922	Jun 09, 2030	DP			
<u>ALENDRONATE SODIUM - BINOSTO</u>						
N 202344 001	9592195	Dec 05, 2031	DP			
<u>ALISKIREN HEMIFUMARATE - TEKTURNA</u>						
N 021985 001	8617595	Feb 19, 2026	DP			
	8617595*PED	Aug 19, 2026				
<u>ALISKIREN HEMIFUMARATE - TEKTURNA</u>						
N 021985 002	8617595	Feb 19, 2026	DP			
	8617595*PED	Aug 19, 2026				

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>ALISKIREN HEMIFUMARATE; AMLODIPINE BESYLATE - TEKAMLO</u>						
N 022545	001	8613949	Dec 21, 2029	DP		
<u>ALISKIREN HEMIFUMARATE; AMLODIPINE BESYLATE - TEKAMLO</u>						
N 022545	002	8613949	Dec 21, 2029	DP		
<u>ALISKIREN HEMIFUMARATE; AMLODIPINE BESYLATE - TEKAMLO</u>						
N 022545	003	8613949	Dec 21, 2029	DP		
<u>ALISKIREN HEMIFUMARATE; AMLODIPINE BESYLATE - TEKAMLO</u>						
N 022545	004	8613949	Dec 21, 2029	DP		
<u>ALISKIREN HEMIFUMARATE; HYDROCHLOROTHIAZIDE - TEKTURNA HCT</u>						
N 022107	001	8618172	Jul 13, 2028	DP		
<u>ALISKIREN HEMIFUMARATE; HYDROCHLOROTHIAZIDE - TEKTURNA HCT</u>						
N 022107	002	8618172	Jul 13, 2028	DP		
<u>ALISKIREN HEMIFUMARATE; HYDROCHLOROTHIAZIDE - TEKTURNA HCT</u>						
N 022107	003	8618172	Jul 13, 2028	DP		
<u>ALISKIREN HEMIFUMARATE; HYDROCHLOROTHIAZIDE - TEKTURNA HCT</u>						
N 022107	004	8618172	Jul 13, 2028	DP		
<u>ALISKIREN HEMIFUMARATE; VALSARTAN - VALTURNA</u>						
N 022217	001	8168616	Jul 03, 2026	DP		
<u>ALISKIREN HEMIFUMARATE; VALSARTAN - VALTURNA</u>						
N 022217	002	8168616	Jul 03, 2026	DP		
<u>ALLOPURINOL; LESINURAD - DUZALLO</u>						
N 209203	001	10183012	Nov 26, 2028			U-2104
		8003681	Aug 25, 2025	DS		
		8084483	Aug 17, 2029			U-2104
		8283369	Nov 26, 2028			U-2104
		8357713	Dec 22, 2029		DP	U-2104
		8546436	Feb 29, 2032	DS		
		8546437	Apr 29, 2029			U-2104
		9216179	Aug 01, 2031			U-2104
		9956205	Dec 28, 2031			U-2104
<u>ALLOPURINOL; LESINURAD - DUZALLO</u>						
N 209203	002	10183012	Nov 26, 2028			U-2104
		8003681	Aug 25, 2025	DS		
		8084483	Aug 17, 2029			U-2104
		8283369	Nov 26, 2028			U-2104
		8357713	Dec 22, 2029		DP	U-2104
		8546436	Feb 29, 2032	DS		
		8546437	Apr 29, 2029			U-2104
		9216179	Aug 01, 2031			U-2104
		9956205	Dec 28, 2031			U-2104
<u>ALOGLIPTIN BENZOATE - NESINA</u>						
N 022271	001	7807689	Jun 27, 2028	DS DP	U-1337	
		8173663	Dec 02, 2025		U-1338	
		8288539	Mar 15, 2025	DS		
		8697125	Jun 16, 2029	DP		
					M-300	Jul 27, 2026

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>ALOGLIPTIN BENZOATE - NESINA</u>						
N 022271 002	7807689	Jun 27, 2028	DS DP U-1337		M-300	Jul 27, 2026
	8173663	Dec 02, 2025	U-1338			
	8288539	Mar 15, 2025	DS			
	8697125	Jun 16, 2029	DP			
<u>ALOGLIPTIN BENZOATE - NESINA</u>						
N 022271 003	7807689	Jun 27, 2028	DS DP U-1337		M-300	Jul 27, 2026
	8173663	Dec 02, 2025	U-1338			
	8288539	Mar 15, 2025	DS			
	8697125	Jun 16, 2029	DP			
<u>ALOGLIPTIN BENZOATE; METFORMIN HYDROCHLORIDE - KAZANO</u>						
N 203414 001	7807689	Jun 27, 2028	DS DP U-1337		M-300	Jul 27, 2026
	8173663	Mar 15, 2025	U-1338			
	8288539	Jun 24, 2025	DS			
	8900638	May 24, 2029	DP			
<u>ALOGLIPTIN BENZOATE; METFORMIN HYDROCHLORIDE - KAZANO</u>						
N 203414 002	7807689	Jun 27, 2028	DS DP U-1337		M-300	Jul 27, 2026
	8173663	Mar 15, 2025	U-1338			
	8288539	Jun 24, 2025	DS			
	8900638	May 24, 2029	DP			
<u>ALOGLIPTIN BENZOATE; PIOGLITAZONE HYDROCHLORIDE - OSENI</u>						
N 022426 001	7807689	Jun 27, 2028	DS DP U-1337			
	8173663	Mar 15, 2025	U-1338			
	8288539	Mar 15, 2025	DS			
	8637079	Jun 04, 2029	DP			
<u>ALOGLIPTIN BENZOATE; PIOGLITAZONE HYDROCHLORIDE - OSENI</u>						
N 022426 002	7807689	Jun 27, 2028	DS DP U-1337			
	8173663	Mar 15, 2025	U-1338			
	8288539	Mar 15, 2025	DS			
	8637079	Jun 04, 2029	DP			
<u>ALOGLIPTIN BENZOATE; PIOGLITAZONE HYDROCHLORIDE - OSENI</u>						
N 022426 003	7807689	Jun 27, 2028	DS DP U-1337			
	8173663	Mar 15, 2025	U-1338			
	8288539	Mar 15, 2025	DS			
	8637079	Jun 04, 2029	DP			
<u>ALOGLIPTIN BENZOATE; PIOGLITAZONE HYDROCHLORIDE - OSENI</u>						
N 022426 004	7807689	Jun 27, 2028	DS DP U-1337			
	8173663	Mar 15, 2025	U-1338			
	8288539	Mar 15, 2025	DS			
	8637079	Jun 04, 2029	DP			
<u>ALOGLIPTIN BENZOATE; PIOGLITAZONE HYDROCHLORIDE - OSENI</u>						
N 022426 005	7807689	Jun 27, 2028	DS DP U-1337			
	8173663	Mar 15, 2025	U-1338			
	8288539	Mar 15, 2025	DS			
	8637079	Jun 04, 2029	DP			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>ALOGLIPTIN BENZOATE; PIOGLITAZONE HYDROCHLORIDE - OSENI</u>						
N 022426	006 7807689	Jun 27, 2028	DS DP U-1337			
	8173663	Mar 15, 2025	U-1338			
	8288539	Mar 15, 2025	DS			
	8637079	Jun 04, 2029	DP			
<u>ALPELISIB - PIORAY</u>						
N 212526	001 8227462	Apr 29, 2033	DS DP U-3809		I-937	Jan 18, 2027
	8476268	Sep 10, 2029	DS DP			
<u>ALPELISIB - PIORAY</u>						
N 212526	002 8227462	Apr 29, 2033	DS DP U-3809		I-937	Jan 18, 2027
	8476268	Sep 10, 2029	DS DP			
<u>ALPELISIB - PIORAY</u>						
N 212526	003 8227462	Apr 29, 2033	DS DP U-3809		I-937	Jan 18, 2027
	8476268	Sep 10, 2029	DS DP			
<u>ALPELISIB - VIJOICE</u>						
N 215039	001 8227462	Apr 29, 2033	DS DP		ODE-396	Apr 05, 2029
	8476268	Sep 10, 2029	DS DP			
<u>ALPELISIB - VIJOICE</u>						
N 215039	002 8227462	Apr 29, 2033	DS DP		ODE-396	Apr 05, 2029
	8476268	Sep 10, 2029	DS DP			
<u>ALPELISIB - VIJOICE</u>						
N 215039	003 8227462	Apr 29, 2033	DS DP		ODE-396	Apr 05, 2029
	8476268	Sep 10, 2029	DS DP			
<u>ALPELISIB - VIJOICE</u>						
N 218466	001 8227462	Apr 29, 2033	DS DP			
	8476268	Sep 10, 2029	DS DP			
<u>ALVIMOPAN - ENTEREG</u>						
N 021775	001 8946262	Feb 12, 2030	U-1655			
<u>AMANTADINE HYDROCHLORIDE - GOCOVRI</u>						
N 208944	001 10154971	Dec 04, 2034	U-2459			
	10646456	Jun 17, 2034	U-2808			
	11065213	Aug 23, 2038	DP			
	11077073	Aug 23, 2038	U-2106			
	11077073	Aug 23, 2038	U-2224			
	11077073	Aug 23, 2038	U-3180			
	11197835	Dec 02, 2030	U-2106			
	11903908	Jun 17, 2034	U-3822			
	8389578	Jan 22, 2028	U-2105			
	8741343	Dec 02, 2030	U-2106			
	8796337	Nov 23, 2025	U-2106			
	8889740	Nov 23, 2025	DP			
	8895614	Nov 23, 2025	DP			
	8895615	Nov 23, 2025	U-2106			
	8895616	Nov 23, 2025	U-2106			
	8895617	Nov 23, 2025	U-2106			
	8895618	Nov 23, 2025	DP			
	9867791	Dec 02, 2030	U-2106			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>AMANTADINE HYDROCHLORIDE - GOCOVRI</u>						
N 208944	001	9867792	Dec 02, 2030			U-2106
		9867793	Dec 02, 2030			U-2106
		9877933	Dec 02, 2030			U-2224
<u>AMANTADINE HYDROCHLORIDE - GOCOVRI</u>						
N 208944	002	10154971	Dec 04, 2034			U-2459
		10646456	Jun 17, 2034			U-2808
		11065213	Aug 23, 2038	DP		
		11077073	Aug 23, 2038			U-2106
		11077073	Aug 23, 2038			U-2224
		11077073	Aug 23, 2038			U-3180
		11197835	Dec 02, 2030			U-2106
		11903908	Jun 17, 2034			U-3822
		8389578	Jan 22, 2028			U-2105
		8741343	Dec 02, 2030			U-2106
		8796337	Nov 23, 2025			U-2106
		8889740	Nov 23, 2025	DP		
		8895614	Nov 23, 2025	DP		
		8895615	Nov 23, 2025			U-2106
		8895616	Nov 23, 2025			U-2106
		8895617	Nov 23, 2025			U-2106
		8895618	Nov 23, 2025	DP		
		9867791	Dec 02, 2030			U-2106
		9867792	Dec 02, 2030			U-2106
		9867793	Dec 02, 2030			U-2106
		9877933	Dec 02, 2030			U-2224
<u>AMANTADINE HYDROCHLORIDE - OSMOLEX ER</u>						
N 209410	001	10213393	Feb 15, 2038			U-20
		10213394	Feb 15, 2038			U-2497
		10500170	Feb 15, 2038			U-20
		10500171	Feb 15, 2038			U-2497
		10500172	Feb 15, 2038			U-2497
		10512617	Feb 15, 2038			U-2497
		11890261	Feb 15, 2038			U-2497
		8252331	Mar 13, 2030	DP		
		8389578	Jan 22, 2028			U-219
		8389578	Jan 22, 2028			U-3054
		8574626	Nov 28, 2025	DP		U-20
		8796337	Nov 23, 2025			U-219
		8796337	Nov 23, 2025			U-2497
		8796337	Nov 23, 2025			U-3054
		8889740	Nov 23, 2025	DP		
		8895614	Nov 23, 2025	DP		
		8895615	Nov 23, 2025			U-219
		8895615	Nov 23, 2025			U-3054
		8895616	Nov 23, 2025			U-219
		8895616	Nov 23, 2025			U-3054
		8895617	Nov 23, 2025			U-219
		8895617	Nov 23, 2025			U-3054
		8895618	Nov 23, 2025	DP		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>AMANTADINE HYDROCHLORIDE - OSMOLEX ER</u>						
N 209410 001	8987333	Nov 23, 2025	DP			
	9072697	Nov 23, 2025		U-219		
	9072697	Nov 23, 2025		U-3054		
<u>AMANTADINE HYDROCHLORIDE - OSMOLEX ER</u>						
N 209410 002	10213393	Feb 15, 2038		U-20		
	10213394	Feb 15, 2038		U-2497		
	10500170	Feb 15, 2038		U-20		
	10500171	Feb 15, 2038		U-2497		
	10500172	Feb 15, 2038		U-2497		
	10512617	Feb 15, 2038		U-2497		
	11890261	Feb 15, 2038		U-2497		
	8252331	Mar 13, 2030	DP			
	8389578	Jan 22, 2028		U-219		
	8389578	Jan 22, 2028		U-3054		
	8574626	Nov 28, 2025	DP	U-20		
	8796337	Nov 23, 2025		U-219		
	8796337	Nov 23, 2025		U-2497		
	8796337	Nov 23, 2025		U-3054		
	8889740	Nov 23, 2025	DP			
	8895614	Nov 23, 2025	DP			
	8895615	Nov 23, 2025		U-219		
	8895615	Nov 23, 2025		U-3054		
	8895616	Nov 23, 2025		U-219		
	8895616	Nov 23, 2025		U-3054		
	8895617	Nov 23, 2025		U-219		
	8895617	Nov 23, 2025		U-3054		
	8895618	Nov 23, 2025	DP			
	8987333	Nov 23, 2025	DP			
	9072697	Nov 23, 2025		U-219		
	9072697	Nov 23, 2025		U-3054		
<u>AMANTADINE HYDROCHLORIDE - OSMOLEX ER</u>						
N 209410 003	10213393	Feb 15, 2038		U-20		
	10213394	Feb 15, 2038		U-2497		
	10500170	Feb 15, 2038		U-20		
	10500171	Feb 15, 2038		U-2497		
	10500172	Feb 15, 2038		U-2497		
	10512617	Feb 15, 2038		U-2497		
	8252331	Mar 13, 2030	DP			
	8389578	Jan 22, 2028		U-219		
	8389578	Jan 22, 2028		U-3054		
	8574626	Nov 28, 2025	DP	U-20		
	8796337	Nov 23, 2025		U-219		
	8796337	Nov 23, 2025		U-2497		
	8796337	Nov 23, 2025		U-3054		
	8889740	Nov 23, 2025	DP			
	8895614	Nov 23, 2025	DP			
	8895615	Nov 23, 2025		U-219		
	8895615	Nov 23, 2025		U-3054		
	8895616	Nov 23, 2025		U-219		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>AMANTADINE HYDROCHLORIDE - OSMOLEX ER</u>						
N 209410	003	8895616	Nov 23, 2025		U-3054	
		8895617	Nov 23, 2025		U-219	
		8895617	Nov 23, 2025		U-3054	
		8895618	Nov 23, 2025	DP		
		8987333	Nov 23, 2025	DP		
		9072697	Nov 23, 2025		U-219	
		9072697	Nov 23, 2025		U-3054	
<u>AMANTADINE HYDROCHLORIDE - OSMOLEX ER</u>						
N 209410	004	10213393	Feb 15, 2038		U-20	
		10213394	Feb 15, 2038		U-2497	
		10500170	Feb 15, 2038		U-20	
		10500171	Feb 15, 2038		U-2497	
		10500172	Feb 15, 2038		U-2497	
		10512617	Feb 15, 2038		U-2497	
		8252331	Mar 13, 2030	DP		
		8389578	Jan 22, 2028		U-219	
		8389578	Jan 22, 2028		U-3054	
		8574626	Nov 28, 2025	DP	U-20	
		8796337	Nov 23, 2025		U-219	
		8796337	Nov 23, 2025		U-2497	
		8796337	Nov 23, 2025		U-3054	
		8889740	Nov 23, 2025	DP		
		8895614	Nov 23, 2025	DP		
		8895615	Nov 23, 2025		U-219	
		8895615	Nov 23, 2025		U-3054	
		8895616	Nov 23, 2025		U-219	
		8895616	Nov 23, 2025		U-3054	
		8895617	Nov 23, 2025		U-219	
		8895617	Nov 23, 2025		U-3054	
		8895618	Nov 23, 2025	DP		
		8987333	Nov 23, 2025	DP		
		9072697	Nov 23, 2025		U-219	
		9072697	Nov 23, 2025		U-3054	
<u>AMBRISANTAN - LETAIRIS</u>						
N 022081	001	8377933	Dec 11, 2027		U-1754	
		9474752	Dec 11, 2027		U-1754	
		9549926	Oct 14, 2031		U-1965	
<u>AMBRISANTAN - LETAIRIS</u>						
N 022081	002	8377933	Dec 11, 2027		U-1754	
		9474752	Dec 11, 2027		U-1754	
		9549926	Oct 14, 2031		U-1965	
<u>AMIFAMPRIDINE PHOSPHATE - FIRDAPSE</u>						
N 208078	001	10626088	Feb 25, 2037	DP	NPP	Sep 29, 2025
		10793893	May 26, 2034		U-2956	ODE-223 Nov 28, 2025
		11060128	Jun 29, 2032		U-2956	
		11268128	Jun 29, 2032		U-2956	
		11274331	Jun 29, 2032		U-2956	
		11274332	Jun 29, 2032		U-2956	

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>AMIFAMPRIDINE PHOSPHATE - FIRDAPSE</u>						
N 208078 001	10626088	Feb 25, 2037	DP		NPP	Sep 29, 2025
	10793893	May 26, 2034	U-2956		ODE-223	Nov 28, 2025
	11060128	Jun 29, 2032	U-2956			
	11268128	Jun 29, 2032	U-2956			
	11274331	Jun 29, 2032	U-2956			
	11274332	Jun 29, 2032	U-2956			
<u>AMIKACIN SULFATE - ARIKAYCE KIT</u>						
N 207356 001	10251900	May 15, 2035	U-2414		ODE-214	Sep 28, 2025
	10751355	May 15, 2035	U-2414		GAIN	Sep 28, 2030
	11446318	May 15, 2035	U-2414			
	12016873	May 15, 2035	U-2414			
	7718189	Jun 06, 2025	DP U-2415			
	8226975	Aug 15, 2028	DP			
	8632804	Dec 05, 2026	U-2416			
	8642075	Dec 05, 2026	DP			
	8679532	Dec 05, 2026	U-2415			
	9566234	Jan 18, 2034	DP U-2415			
	9895385	May 15, 2035	U-2417			
<u>AMINOLEVULINIC ACID HYDROCHLORIDE - LEVULAN</u>						
N 020965 001	10357567	Jan 12, 2038	U-3163			
	11077192	Jan 12, 2038	U-3163			
	11135293	Jan 12, 2038	U-3163			
	11571478	Jan 12, 2038	U-3163			
	11690914	Jan 12, 2038	U-3163			
<u>AMINOLEVULINIC ACID HYDROCHLORIDE - AMELUZ</u>						
N 208081 001	11235169	Oct 15, 2040	U-3303		D-194	Oct 04, 2027
	11540981	Feb 07, 2028	DP			
<u>AMIODARONE HYDROCHLORIDE - NEXTERONE</u>						
N 022325 001	7635773	Mar 13, 2029	DP			
<u>AMIODARONE HYDROCHLORIDE - NEXTERONE</u>						
N 022325 002	7635773	Mar 13, 2029	DP			
<u>AMIODARONE HYDROCHLORIDE - NEXTERONE</u>						
N 022325 003	7635773	Mar 13, 2029	DP			
<u>AMISULPRIDE - BARHEMSYS</u>						
N 209510 001	10525033	Mar 10, 2031	DP		NCE	Feb 26, 2025
	11357753	Feb 09, 2038	U-2754			
	9084765	Feb 26, 2034	U-1744			
	9084765	Feb 26, 2034	U-2754			
	9084765	Feb 26, 2034	U-3467			
	9545426	Mar 10, 2031	U-1744			
	9545426	Mar 10, 2031	U-2754			
	9889118	Mar 10, 2031	U-1744			
	9889118	Mar 10, 2031	U-2754			
<u>AMISULPRIDE - BARHEMSYS</u>						
N 209510 002	10525033	Mar 10, 2031	DP		NCE	Feb 26, 2025
	11357753	Feb 09, 2038	U-2754			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>AMISULPRIDE - BARHEMSYS</u>						
N 209510	002	9084765	Feb 26, 2034	U-1744		
		9084765	Feb 26, 2034	U-2754		
		9084765	Feb 26, 2034	U-3467		
		9545426	Mar 10, 2031	U-1744		
		9545426	Mar 10, 2031	U-2754		
		9889118	Mar 10, 2031	U-1744		
		9889118	Mar 10, 2031	U-2754		
<u>AMLODIPINE BENZOATE - KATERZIA</u>						
N 211340	001	10695329	Oct 16, 2037	DP		
		10799453	Apr 11, 2039	DP		
		10894039	Oct 06, 2037	U-185		
		10894039	Oct 06, 2037	U-3		
		10952998	Oct 06, 2037	DP		
		10959991	Oct 06, 2037	U-158		
		10959991	Oct 06, 2037	U-39		
		11364230	Oct 06, 2037	DP		
		11471409	Oct 06, 2037	U-3447		
		11471409	Oct 06, 2037	U-3448		
		11484498	Oct 06, 2037	DP		
		11701326	Oct 06, 2037	DP		
		11918685	Oct 06, 2037	U-3447		
		11918685	Oct 06, 2037	U-3448		
		12053461	Oct 06, 2037	DP		
<u>AMLODIPINE BESYLATE - NORLIOVA</u>						
N 214439	001	11253474	Feb 24, 2041	DP U-3309		
		11253474	Feb 24, 2041	DP U-3310		
		11253474	Feb 24, 2041	DP U-3311		
		11458095	Feb 24, 2041	DP U-3309		
		11458095	Feb 24, 2041	DP U-3310		
		11458095	Feb 24, 2041	DP U-3311		
		11723866	Feb 24, 2041	DP U-3309		
		11723866	Feb 24, 2041	DP U-3310		
		11723866	Feb 24, 2041	DP U-3311		
		12005141	Feb 24, 2041	DP U-3309		
		12005141	Feb 24, 2041	DP U-3310		
<u>AMLODIPINE BESYLATE; CELECOXIB - CONSENSI</u>						
N 210045	001	10350171	Jun 14, 2038	DP		
		10925835	Jun 14, 2038	U-2410		
		10945960	Jun 14, 2038	DP		
		9408837	Feb 28, 2030	U-2410		
		9662315	May 22, 2029	DP U-2410		
<u>AMLODIPINE BESYLATE; CELECOXIB - CONSENSI</u>						
N 210045	002	10350171	Jun 14, 2038	DP		
		10925835	Jun 14, 2038	U-2410		
		10945960	Jun 14, 2038	DP		
		9408837	Feb 28, 2030	U-2410		
		9662315	May 22, 2029	DP U-2410		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>AMLODIPINE BESYLATE; CELECOXIB - CONSENSI</u>						
N 210045	003 10350171	Jun 14, 2038	DP			
	10925835	Jun 14, 2038	U-2410			
	10945960	Jun 14, 2038	DP			
	9408837	Feb 28, 2030	U-2410			
	9662315	May 22, 2029	DP U-2410			
<u>AMLODIPINE BESYLATE; PERINDOPRIL ARGININE - PRESTALIA</u>						
N 205003	001 7846961	Oct 05, 2029	DS DP U-3			
<u>AMLODIPINE BESYLATE; PERINDOPRIL ARGININE - PRESTALIA</u>						
N 205003	002 7846961	Oct 05, 2029	DS DP U-3			
<u>AMLODIPINE BESYLATE; PERINDOPRIL ARGININE - PRESTALIA</u>						
N 205003	003 7846961	Oct 05, 2029	DS DP U-3			
<u>AMOXICILLIN - MOXATAG</u>						
N 050813	001 8299052	May 07, 2027	U-1304			
	8357394	Dec 08, 2026	DP			
	8778924	Dec 08, 2026	DS DP U-897			
<u>AMOXICILLIN; CLARITHROMYCIN; VONOPRAZAN FUMARATE - VOQUEZNA TRIPLE PAK</u>						
N 215152	001 7977488	Aug 11, 2028	DS		NCE	May 03, 2027
	9186411	Aug 11, 2030	DP		GAIN	May 03, 2032
<u>AMOXICILLIN; OMEPRAZOLE MAGNESIUM; RIFABUTIN - TALICIA</u>						
N 213004	001 10238606	Feb 12, 2034	DP		NP	Nov 01, 2022
	11135172	Feb 12, 2034	DP U-2660		GAIN	Nov 01, 2027
	11878011	May 27, 2042	U-3816			
	11931463	Feb 12, 2034	DP U-3866			
	9050263	Feb 12, 2034	DP U-2660			
	9498445	Feb 12, 2034	DP U-2660			
	9603806	Feb 12, 2034	DP U-2660			
<u>AMOXICILLIN; VONOPRAZAN FUMARATE - VOQUEZNA DUAL PAK</u>						
N 215153	001 7977488	Aug 11, 2028	DS		NCE	May 03, 2027
	9186411	Aug 11, 2030	DP		GAIN	May 03, 2032
<u>AMPHETAMINE - ADZENYS ER</u>						
N 204325	001 8709491	Jun 28, 2032	DP			
	9017731	Jun 28, 2032	DP			
	9265737	Jun 28, 2032	DP			
<u>AMPHETAMINE - ADZENYS XR-ODT</u>						
N 204326	001 8709491	Jun 28, 2032	DP			
	8840924	Apr 09, 2026	DP			
	9017731	Jun 28, 2032	DP			
	9265737	Jun 28, 2032	DP			
	9839619	Jun 28, 2032	DP			
<u>AMPHETAMINE - ADZENYS XR-ODT</u>						
N 204326	002 8709491	Jun 28, 2032	DP			
	8840924	Apr 09, 2026	DP			
	9017731	Jun 28, 2032	DP			
	9265737	Jun 28, 2032	DP			
	9839619	Jun 28, 2032	DP			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>AMPHETAMINE - ADZENYS XR-ODT</u>						
N 204326	003	8709491	Jun 28, 2032	DP		
		8840924	Apr 09, 2026	DP		
		9017731	Jun 28, 2032	DP		
		9265737	Jun 28, 2032	DP		
		9839619	Jun 28, 2032	DP		
<u>AMPHETAMINE - ADZENYS XR-ODT</u>						
N 204326	004	8709491	Jun 28, 2032	DP		
		8840924	Apr 09, 2026	DP		
		9017731	Jun 28, 2032	DP		
		9265737	Jun 28, 2032	DP		
		9839619	Jun 28, 2032	DP		
<u>AMPHETAMINE - ADZENYS XR-ODT</u>						
N 204326	005	8709491	Jun 28, 2032	DP		
		8840924	Apr 09, 2026	DP		
		9017731	Jun 28, 2032	DP		
		9265737	Jun 28, 2032	DP		
		9839619	Jun 28, 2032	DP		
<u>AMPHETAMINE - ADZENYS XR-ODT</u>						
N 204326	006	8709491	Jun 28, 2032	DP		
		8840924	Apr 09, 2026	DP		
		9017731	Jun 28, 2032	DP		
		9265737	Jun 28, 2032	DP		
		9839619	Jun 28, 2032	DP		
<u>AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATE - MYDAYIS</u>						
N 022063	001	8846100	Aug 24, 2029	DP		
		9173857	May 12, 2026		U-2025	
<u>AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATE - MYDAYIS</u>						
N 022063	002	8846100	Aug 24, 2029	DP		
		9173857	May 12, 2026		U-2025	
<u>AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATE - MYDAYIS</u>						
N 022063	003	8846100	Aug 24, 2029	DP		
		9173857	May 12, 2026		U-2025	
<u>AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATE - MYDAYIS</u>						
N 022063	004	8846100	Aug 24, 2029	DP		
		9173857	May 12, 2026		U-2025	
<u>AMPHETAMINE SULFATE - EVEKEO ODT</u>						
N 209905	001	10441554	Mar 10, 2037	DP		
		11160772	Mar 10, 2037	DP		
		11896562	Mar 10, 2037		U-3299	
<u>AMPHETAMINE SULFATE - EVEKEO ODT</u>						
N 209905	002	10441554	Mar 10, 2037	DP		
		11160772	Mar 10, 2037	DP		
		11896562	Mar 10, 2037		U-3299	

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>AMPHETAMINE SULFATE - EVEKEO ODT</u>						
N 209905	003	10441554	Mar 10, 2037	DP		
		11160772	Mar 10, 2037	DP		
		11896562	Mar 10, 2037		U-3299	
<u>AMPHETAMINE SULFATE - EVEKEO ODT</u>						
N 209905	004	10441554	Mar 10, 2037	DP		
		11160772	Mar 10, 2037	DP		
		11896562	Mar 10, 2037		U-3299	
<u>AMPHETAMINE SULFATE - EVEKEO ODT</u>						
N 209905	005	10441554	Mar 10, 2037	DP		
		11160772	Mar 10, 2037	DP		
<u>AMPHETAMINE; AMPHETAMINE ASPARTATE/DEXTROAMPHETAMINE SULFATE - DYANA VEL XR</u>						
N 208147	001	10086087	Mar 15, 2027	DP		
		11590228	Sep 07, 2036	DP	U-3538	
		8062667	Mar 29, 2029	DP		
		8597684	Mar 15, 2027	DP		
		8747902	Mar 15, 2027	DP		
		8883217	Mar 15, 2027	DP		
		9675703	Mar 15, 2027	DP		
<u>AMPHETAMINE; AMPHETAMINE ASPARTATE/DEXTROAMPHETAMINE SULFATE - DYANA VEL XR 5</u>						
N 210526	001	11590081	Sep 24, 2038	DP	U-3538	
		8337890	Mar 15, 2027	DP		
		8747902	Mar 15, 2027	DP		
		9675704	Mar 15, 2027	DP		
<u>AMPHETAMINE; AMPHETAMINE ASPARTATE/DEXTROAMPHETAMINE SULFATE - DYANA VEL XR 10</u>						
N 210526	002	11590081	Sep 24, 2038	DP	U-3538	
		8337890	Mar 15, 2027	DP		
		8747902	Mar 15, 2027	DP		
		9675704	Mar 15, 2027	DP		
<u>AMPHETAMINE; AMPHETAMINE ASPARTATE/DEXTROAMPHETAMINE SULFATE - DYANA VEL XR 15</u>						
N 210526	003	11590081	Sep 24, 2038	DP	U-3538	
		8337890	Mar 15, 2027	DP		
		8747902	Mar 15, 2027	DP		
		9675704	Mar 15, 2027	DP		
<u>AMPHETAMINE; AMPHETAMINE ASPARTATE/DEXTROAMPHETAMINE SULFATE - DYANA VEL XR 20</u>						
N 210526	004	11590081	Sep 24, 2038	DP	U-3538	
		8337890	Mar 15, 2027	DP		
		8747902	Mar 15, 2027	DP		
		9675704	Mar 15, 2027	DP		
<u>ANGIOTENSIN II ACETATE - GIAPREZA</u>						
N 209360	001	10028995	Dec 18, 2034		U-2338	
		10335451	Dec 16, 2029		U-2581	
		10493124	Dec 18, 2034		U-2679	
		10500247	Dec 16, 2029		U-2680	
		10500247	Dec 16, 2029		U-2681	
		10548943	Dec 16, 2029		U-2739	
		10548943	Dec 16, 2029		U-2740	

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>ANGIOTENSIN II ACETATE - GIAPREZA</u>						
N 209360 001	11096983	Dec 18, 2034	U-3211			
	11096983	Dec 18, 2034	U-3212			
	11219662	Jan 06, 2037	U-3262			
	11559559	Dec 18, 2034	U-3514			
	9220745	Dec 18, 2034	U-2217			
	9220745	Dec 18, 2034	U-2218			
	9572856	Jul 18, 2031	U-2221			
	9867863	Dec 16, 2029	U-2231			
<u>ANGIOTENSIN II ACETATE - GIAPREZA</u>						
N 209360 002	10028995	Dec 18, 2034	U-2338			
	10335451	Dec 16, 2029	U-2581			
	10493124	Dec 18, 2034	U-2679			
	10500247	Dec 16, 2029	U-2680			
	10500247	Dec 16, 2029	U-2681			
	10548943	Dec 16, 2029	U-2739			
	10548943	Dec 16, 2029	U-2740			
	11096983	Dec 18, 2034	U-3211			
	11096983	Dec 18, 2034	U-3212			
	11219662	Jan 06, 2037	U-3262			
	11559559	Dec 18, 2034	U-3514			
	9220745	Dec 18, 2034	U-2217			
	9220745	Dec 18, 2034	U-2218			
	9572856	Nov 20, 2030	U-2221			
	9867863	Dec 16, 2029	U-2231			
<u>ANGIOTENSIN II ACETATE - GIAPREZA</u>						
N 209360 003	10028995	Dec 18, 2034	U-2338			
	10335451	Dec 16, 2029	U-2581			
	10493124	Dec 18, 2034	U-2679			
	10500247	Dec 16, 2029	U-2680			
	10500247	Dec 16, 2029	U-2681			
	10548943	Dec 16, 2029	U-2739			
	10548943	Dec 16, 2029	U-2740			
	11096983	Dec 18, 2034	U-3211			
	11096983	Dec 18, 2034	U-3212			
	11219662	Jan 06, 2037	U-3262			
	11559559	Dec 18, 2034	U-3514			
	9220745	Dec 18, 2034	U-2217			
	9220745	Dec 18, 2034	U-2218			
	9572856	Jul 18, 2031	U-2221			
	9867863	Dec 16, 2029	U-2231			
<u>APALUTAMIDE - ERLEADA</u>						
N 210951 001	10052314	Sep 23, 2033	U-2381		Y	
	10052314	Sep 23, 2033	U-2382		Y	
	10702508	Apr 30, 2038	U-3012			
	10849888	Sep 23, 2033	U-3013			
	11963952	Jan 30, 2040	U-3901			
	8445507	Sep 15, 2030	DS DP U-2237			
	8445507	Sep 15, 2030	DS DP U-2624			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>APALUTAMIDE - ERLEADA</u>						
N 210951	001	8802689	Mar 27, 2027		U-2237	
		8802689	Mar 27, 2027		U-2624	
		9388159	Mar 27, 2027	DS DP		
		9481663	Jun 04, 2033	DS DP	U-2237	
		9481663	Jun 04, 2033	DS DP	U-2624	
		9884054	Sep 23, 2033		U-2237	
		9987261	Mar 27, 2027	DP		
		RE49353	Sep 23, 2033		U-2381	
<u>APALUTAMIDE - ERLEADA</u>						
N 210951	002	10702508	Apr 30, 2038		U-3012	
		10849888	Sep 23, 2033		U-3013	
		11963952	Jan 30, 2040		U-3901	
		8445507	Sep 15, 2030	DS DP	U-2237	
		8445507	Sep 15, 2030	DS DP	U-2624	
		8802689	Mar 27, 2027		U-2237	
		8802689	Mar 27, 2027		U-2624	
		9388159	Mar 27, 2027	DS DP		
		9481663	Jun 04, 2033	DS DP	U-2237	
		9481663	Jun 04, 2033	DS DP	U-2624	
		9884054	Sep 23, 2033		U-2237	
		9987261	Mar 27, 2027	DP		
		RE49353	Sep 23, 2033		U-2381	
<u>APIXABAN - ELIQUIS</u>						
N 202155	001	6967208	Nov 21, 2026	DS DP	U-1167	
		6967208	Nov 21, 2026	DS DP	U-1200	
		6967208	Nov 21, 2026	DS DP	U-1301	
		6967208	Nov 21, 2026	DS DP	U-1302	
		6967208	Nov 21, 2026	DS DP	U-1323	
		6967208	Nov 21, 2026	DS DP	U-1501	
		6967208	Nov 21, 2026	DS DP	U-1502	
		6967208	Nov 21, 2026	DS DP	U-1729	
		6967208	Nov 21, 2026	DS DP	U-1730	
		9326945	Feb 24, 2031	DP		
<u>APIXABAN - ELIQUIS</u>						
N 202155	002	6967208	Nov 21, 2026	DS DP	U-1200	
		6967208	Nov 21, 2026	DS DP	U-1301	
		6967208	Nov 21, 2026	DS DP	U-1302	
		6967208	Nov 21, 2026	DS DP	U-1323	
		9326945	Feb 24, 2031	DP		
<u>APOMORPHINE HYDROCHLORIDE - KYNMOBI</u>						
N 210875	001	10420763	Jun 11, 2030	DP	U-2825	
		10449146	Apr 19, 2036		U-2825	
		10821074	Aug 07, 2029	DP		
		10959943	Apr 19, 2036		U-2825	
		11419769	Dec 16, 2031	DP	U-2825	
		8414922	Dec 16, 2031	DP	U-2825	
		8846074	Dec 16, 2031	DP	U-2825	
		9044475	Jun 11, 2030	DP		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>APOMORPHINE HYDROCHLORIDE - KYNMOBI</u>						
N 210875	001	9283219	Jun 11, 2030	DP	U-2825	
		9326981	Jun 11, 2030		U-2825	
		9669019	Jun 11, 2030	DP	U-2825	
		9669021	Jun 11, 2030		U-2825	
<u>APOMORPHINE HYDROCHLORIDE - KYNMOBI</u>						
N 210875	002	10420763	Jun 11, 2030	DP	U-2825	
		10449146	Apr 19, 2036		U-2825	
		10821074	Aug 07, 2029	DP		
		10959943	Apr 19, 2036		U-2825	
		11419769	Dec 16, 2031	DP	U-2825	
		8414922	Dec 16, 2031	DP	U-2825	
		8846074	Dec 16, 2031	DP	U-2825	
		9044475	Jun 11, 2030	DP		
		9283219	Jun 11, 2030	DP	U-2825	
		9326981	Jun 11, 2030		U-2825	
		9669019	Jun 11, 2030	DP	U-2825	
		9669021	Jun 11, 2030		U-2825	
<u>APOMORPHINE HYDROCHLORIDE - KYNMOBI</u>						
N 210875	003	10420763	Jun 11, 2030	DP	U-2825	
		10449146	Apr 19, 2036		U-2825	
		10821074	Aug 07, 2029	DP		
		10959943	Apr 19, 2036		U-2825	
		11419769	Dec 16, 2031	DP	U-2825	
		8414922	Dec 16, 2031	DP	U-2825	
		8846074	Dec 16, 2031	DP	U-2825	
		9044475	Jun 11, 2030	DP		
		9283219	Jun 11, 2030	DP	U-2825	
		9326981	Jun 11, 2030		U-2825	
		9669019	Jun 11, 2030	DP	U-2825	
		9669021	Jun 11, 2030		U-2825	
<u>APOMORPHINE HYDROCHLORIDE - KYNMOBI</u>						
N 210875	004	10420763	Jun 11, 2030	DP	U-2825	
		10449146	Apr 19, 2036		U-2825	
		10821074	Aug 07, 2029	DP		
		10959943	Apr 19, 2036		U-2825	
		11419769	Dec 16, 2031	DP	U-2825	
		8414922	Dec 16, 2031	DP	U-2825	
		8846074	Dec 16, 2031	DP	U-2825	
		9044475	Jun 11, 2030	DP		
		9283219	Jun 11, 2030	DP	U-2825	
		9326981	Jun 11, 2030		U-2825	
		9669019	Jun 11, 2030	DP	U-2825	
		9669021	Jun 11, 2030		U-2825	
<u>APOMORPHINE HYDROCHLORIDE - KYNMOBI</u>						
N 210875	005	10420763	Jun 11, 2030	DP	U-2825	
		10449146	Apr 19, 2036		U-2825	
		10821074	Aug 07, 2029	DP		
		10959943	Apr 19, 2036		U-2825	

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>APOMORPHINE HYDROCHLORIDE - KYNMOBI</u>						
N 210875	005	11419769	Dec 16, 2031	DP U-2825		
		8414922	Dec 16, 2031	DP U-2825		
		8846074	Dec 16, 2031	DP U-2825		
		9044475	Jun 11, 2030	DP		
		9283219	Jun 11, 2030	DP U-2825		
		9326981	Jun 11, 2030	U-2825		
		9669019	Jun 11, 2030	DP U-2825		
		9669021	Jun 11, 2030	U-2825		
<u>APREMILAST - OTEZLA</u>						
N 205437	001	10092541	May 29, 2034	U-2403	M-299	Jul 20, 2026
		10092541	May 29, 2034	U-2659	NPP	Apr 25, 2027
		10092541*PED	Nov 29, 2034		ODE-248	Jul 19, 2026
		7427638	Feb 16, 2028	DS DP	PED	Jan 19, 2027
		7427638*PED	Aug 16, 2028		PED	Jan 20, 2027
		9872854	May 29, 2034	U-2232	PED	Oct 25, 2027
		9872854	May 29, 2034	U-2233		
		9872854*PED	Nov 29, 2034			
<u>APREMILAST - OTEZLA</u>						
N 205437	002	10092541	May 29, 2034	U-2403	M-299	Jul 20, 2026
		10092541	May 29, 2034	U-2659	NPP	Apr 25, 2027
		10092541*PED	Nov 29, 2034		ODE-248	Jul 19, 2026
		7427638	Feb 16, 2028	DS DP	PED	Jan 19, 2027
		7427638*PED	Aug 16, 2028		PED	Jan 20, 2027
		9872854	May 29, 2034	U-2232	PED	Oct 25, 2027
		9872854	May 29, 2034	U-2233		
		9872854*PED	Nov 29, 2034			
<u>APREMILAST - OTEZLA</u>						
N 205437	003	10092541	May 29, 2034	U-2403	M-299	Jul 20, 2026
		10092541	May 29, 2034	U-2659	NPP	Apr 25, 2027
		10092541*PED	Nov 29, 2034		ODE-248	Jul 19, 2026
		7427638	Feb 16, 2028	DS DP	PED	Jan 19, 2027
		7427638*PED	Aug 16, 2028		PED	Jan 20, 2027
		9872854	May 29, 2034	U-2232	PED	Oct 25, 2027
		9872854	May 29, 2034	U-2233		
		9872854*PED	Nov 29, 2034			
<u>APREPITANT - EMEND</u>						
N 021549	001	8258132	Sep 26, 2027	DP U-1743		
		8258132	Sep 26, 2027	DP U-901		
<u>APREPITANT - EMEND</u>						
N 021549	002	8258132	Sep 26, 2027	DP U-1743		
		8258132	Sep 26, 2027	DP U-901		
<u>APREPITANT - EMEND</u>						
N 021549	003	8258132	Sep 26, 2027	DP U-1743		
		8258132	Sep 26, 2027	DP U-901		
<u>APREPITANT - EMEND</u>						
N 207865	001	8258132	Sep 26, 2027	DP U-1916		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>APREPITANT - CINVANTI</u>						
N 209296	001	10500208	Sep 18, 2035	DP		
		10624850	Sep 18, 2035	U-2161		
		10953018	Sep 18, 2035	U-2161		
		11173118	Sep 18, 2035	DP		
		11744800	Sep 18, 2035	DP		
		12115254	Sep 18, 2035	DP		
		12115255	Sep 18, 2035	DP		
		9561229	Sep 18, 2035	DP U-2161		
		9808465	Sep 18, 2035	U-2161		
		9974742	Sep 18, 2035	DP		
		9974793	Sep 18, 2035	DP		
		9974794	Sep 18, 2035	DP U-2161		
<u>APREPITANT - APONVIE</u>						
N 216457	001	10500208	Sep 18, 2035	DP		
		10624850	Sep 18, 2035	U-3440		
		10953018	Sep 18, 2035	U-3440		
		11173118	Sep 18, 2035	DP		
		11744800	Sep 18, 2035	DP U-3690		
		11878074	Sep 18, 2035	U-3787		
		12115254	Sep 18, 2035	DP		
		12115255	Sep 18, 2035	DP		
		9561229	Sep 18, 2035	DP U-3440		
		9808465	Sep 18, 2035	U-3440		
		9974742	Sep 18, 2035	DP		
		9974793	Sep 18, 2035	DP		
		9974794	Sep 18, 2035	DP U-3440		
<u>APROCITENTAN - TRYVIO</u>						
N 217686	001	10919881	Feb 26, 2038	DS DP	NCE	Mar 22, 2029
		11174247	Nov 06, 2037	U-3879		
		11680058	Jul 26, 2038	U-3878		
		11787782	Mar 02, 2038	U-3877		
		8324232	Sep 21, 2029	DS DP U-3878		
<u>ARGATROBAN - ARGATROBAN IN SODIUM CHLORIDE</u>						
N 022434	001	7589106	Sep 26, 2027	DP U-1163		
		7687516	Sep 26, 2027	DP U-1164		
<u>ARIMOCLOMOL CITRATE - MIPLYFFA</u>						
N 214927	001	11045460	Aug 19, 2029	U-4021	NCE	Sep 20, 2029
		9289472	Aug 11, 2029	U-4021	ODE-496	Sep 20, 2031
		9884058	Jun 26, 2029	U-4021		
<u>ARIMOCLOMOL CITRATE - MIPLYFFA</u>						
N 214927	002	11045460	Aug 19, 2029	U-4021	NCE	Sep 20, 2029
		9289472	Aug 11, 2029	U-4021	ODE-496	Sep 20, 2031
		9884058	Jun 26, 2029	U-4021		
<u>ARIMOCLOMOL CITRATE - MIPLYFFA</u>						
N 214927	003	11045460	Aug 19, 2029	U-4021	NCE	Sep 20, 2029
		9289472	Aug 11, 2029	U-4021	ODE-496	Sep 20, 2031
		9884058	Jun 26, 2029	U-4021		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>ARIMOCLOMOL CITRATE - MIPLYFFA</u>						
N 214927 004	11045460	Aug 19, 2029	U-4021		NCE	Sep 20, 2029
	9289472	Aug 11, 2029	U-4021		ODE-496	Sep 20, 2031
	9884058	Jun 26, 2029	U-4021			
<u>ARIPIPRAZOLE - ABILIFY</u>						
N 021436 001	8759350	Mar 02, 2027	U-1529			
	9125939	Jul 28, 2026	U-1749			
<u>ARIPIPRAZOLE - ABILIFY</u>						
N 021436 002	8759350	Mar 02, 2027	U-1529			
	9125939	Jul 28, 2026	U-1749			
<u>ARIPIPRAZOLE - ABILIFY</u>						
N 021436 003	8759350	Mar 02, 2027	U-1529			
	9125939	Jul 28, 2026	U-1749			
<u>ARIPIPRAZOLE - ABILIFY</u>						
N 021436 004	8759350	Mar 02, 2027	U-1529			
	9125939	Jul 28, 2026	U-1749			
<u>ARIPIPRAZOLE - ABILIFY</u>						
N 021436 005	8759350	Mar 02, 2027	U-1529			
	9125939	Jul 28, 2026	U-1749			
<u>ARIPIPRAZOLE - ABILIFY</u>						
N 021436 006	8759350	Mar 02, 2027	U-1529			
	9125939	Jul 28, 2026	U-1749			
<u>ARIPIPRAZOLE - ABILIFY</u>						
N 021713 001	8759350	Mar 02, 2027	U-1529			
<u>ARIPIPRAZOLE - ABILIFY</u>						
N 021729 002	8759350	Mar 02, 2027	U-1529			
	9125939	Jul 28, 2026	U-1749			
<u>ARIPIPRAZOLE - ABILIFY</u>						
N 021729 003	8759350	Mar 02, 2027	U-1529			
	9125939	Jul 28, 2026	U-1749			
<u>ARIPIPRAZOLE - ABILIFY</u>						
N 021866 001	7115587*PED	Jan 21, 2025				
<u>ARIPIPRAZOLE - ABILIFY MAINTENA KIT</u>						
N 202971 001	10525057	Mar 08, 2034	U-1632			
	10525057	Mar 08, 2034	U-2723			
	10525057	Mar 08, 2034	U-543			
	10980803	Sep 24, 2033	U-1632			
	10980803	Sep 24, 2033	U-543			
	11154553	Sep 24, 2033	U-1632			
	11154553	Sep 24, 2033	U-3245			
	11154553	Sep 24, 2033	U-814			
	11344547	Sep 24, 2033	U-1632			
	11344547	Sep 24, 2033	U-3245			
	11344547	Sep 24, 2033	U-814			
	11400087	Sep 24, 2033	U-1632			
	11400087	Sep 24, 2033	U-3245			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>ARIPIRAZOLE - ABILIFY MAINTENA KIT</u>						
N 202971 001	11400087	Sep 24, 2033	U-814			
	11648347	Apr 06, 2034	DP			
	8338427	Mar 15, 2025	DP U-1633			
	8338427	Mar 15, 2025	DP U-543			
	8399469	Jun 29, 2025	DS			
<u>ARIPIRAZOLE - ABILIFY MAINTENA KIT</u>						
N 202971 002	10525057	Mar 08, 2034	U-1632			
	10525057	Mar 08, 2034	U-2723			
	10525057	Mar 08, 2034	U-543			
	10980803	Sep 24, 2033	U-1632			
	10980803	Sep 24, 2033	U-543			
	11154553	Sep 24, 2033	U-1632			
	11154553	Sep 24, 2033	U-3245			
	11154553	Sep 24, 2033	U-814			
	11344547	Sep 24, 2033	U-1632			
	11344547	Sep 24, 2033	U-3245			
	11344547	Sep 24, 2033	U-814			
	11400087	Sep 24, 2033	U-1632			
	11400087	Sep 24, 2033	U-3245			
	11400087	Sep 24, 2033	U-814			
	11648347	Apr 06, 2034	DP			
	8338427	Mar 15, 2025	DP U-1633			
	8338427	Mar 15, 2025	DP U-543			
	8399469	Jun 29, 2025	DS			
<u>ARIPIRAZOLE - ABILIFY MAINTENA KIT</u>						
N 202971 003	10525057	Mar 08, 2034	U-1632			
	10525057	Mar 08, 2034	U-2723			
	10525057	Mar 08, 2034	U-543			
	10980803	Sep 24, 2033	U-1632			
	10980803	Sep 24, 2033	U-543			
	11154553	Sep 24, 2033	U-1632			
	11154553	Sep 24, 2033	U-3245			
	11154553	Sep 24, 2033	U-814			
	11344547	Sep 24, 2033	U-1632			
	11344547	Sep 24, 2033	U-3245			
	11344547	Sep 24, 2033	U-814			
	11400087	Sep 24, 2033	U-1632			
	11400087	Sep 24, 2033	U-3245			
	11400087	Sep 24, 2033	U-814			
	11648347	Apr 06, 2034	DP			
	8338427	Mar 15, 2025	DP U-1633			
	8338427	Mar 15, 2025	DP U-543			
	8399469	Jun 29, 2025	DS			
<u>ARIPIRAZOLE - ABILIFY MAINTENA KIT</u>						
N 202971 004	10525057	Mar 08, 2034	U-1632			
	10525057	Mar 08, 2034	U-2723			
	10525057	Mar 08, 2034	U-543			
	10980803	Sep 24, 2033	U-1632			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>ARIPIRAZOLE - ABILIFY MAINTENA KIT</u>						
N 202971 004	10980803	Sep 24, 2033	U-543			
	11154553	Sep 24, 2033	U-1632			
	11154553	Sep 24, 2033	U-3245			
	11154553	Sep 24, 2033	U-814			
	11344547	Sep 24, 2033	U-1632			
	11344547	Sep 24, 2033	U-3245			
	11344547	Sep 24, 2033	U-814			
	11400087	Sep 24, 2033	U-1632			
	11400087	Sep 24, 2033	U-3245			
	11400087	Sep 24, 2033	U-814			
	11648347	Apr 06, 2034	DP			
	8338427	Mar 15, 2025	DP U-1633			
	8338427	Mar 15, 2025	DP U-543			
	8399469	Jun 29, 2025	DS			
<u>ARIPIRAZOLE - ABILIFY MYCITE KIT</u>						
N 207202 001	10441194	Jul 26, 2029	DP			
	10517507	Jun 13, 2032	DP			
	11229378	Jul 11, 2031	DP			
	11464423	Sep 15, 2030	DP			
	11476952	Apr 28, 2026	DP			
	7978064	Sep 14, 2026	DP			
	8114021	Jun 21, 2030	DP			
	8258962	Nov 25, 2030	DP			
	8545402	Apr 27, 2030	DP			
	8547248	Dec 18, 2030	DP U-2167			
	8674825	Apr 09, 2029	DP U-2170			
	8718193	Dec 05, 2029	DP			
	8759350	Mar 02, 2027	U-1529			
	8847766	Mar 29, 2030	DP U-2167			
	8945005	Aug 19, 2029	DP U-2167			
	8956288	Jul 06, 2029	DP U-2167			
	8961412	Nov 17, 2030	DP			
	9060708	Mar 05, 2029	DP			
	9119554	Dec 16, 2028	DP			
	9125939	Jul 28, 2026	U-1749			
	9149577	Dec 15, 2029	DP			
	9258035	Mar 05, 2029	DP			
	9268909	Oct 15, 2033	DP U-2168			
	9320455	Dec 15, 2031	DP			
	9433371	Sep 15, 2029	DP			
	9444503	Nov 19, 2027	DP U-2169			
	9941931	Nov 04, 2030	DP			
<u>ARIPIRAZOLE - ABILIFY MYCITE KIT</u>						
N 207202 002	10441194	Jul 26, 2029	DP			
	10517507	Jun 13, 2032	DP			
	11229378	Jul 11, 2031	DP			
	11464423	Sep 15, 2030	DP			
	11476952	Apr 28, 2026	DP			
	7978064	Sep 14, 2026	DP			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>ARIPIRAZOLE - ABILIFY MYCITE KIT</u>						
N 207202 002	8114021	Jun 21, 2030	DP			
	8258962	Nov 25, 2030	DP			
	8545402	Apr 27, 2030	DP			
	8547248	Dec 18, 2030	DP U-2167			
	8674825	Apr 09, 2029	DP U-2170			
	8718193	Dec 05, 2029	DP			
	8759350	Mar 02, 2027	U-1529			
	8847766	Mar 29, 2030	DP U-2167			
	8945005	Aug 19, 2029	DP U-2167			
	8956288	Jul 06, 2029	DP U-2167			
	8961412	Nov 17, 2030	DP			
	9060708	Mar 05, 2029	DP			
	9119554	Dec 16, 2028	DP			
	9125939	Jul 28, 2026	U-1749			
	9149577	Dec 15, 2029	DP			
	9258035	Mar 05, 2029	DP			
	9268909	Oct 15, 2033	DP U-2168			
	9320455	Dec 15, 2031	DP			
	9433371	Sep 15, 2029	DP			
	9444503	Nov 19, 2027	DP U-2169			
	9941931	Nov 04, 2030	DP			
<u>ARIPIRAZOLE - ABILIFY MYCITE KIT</u>						
N 207202 003	10441194	Jul 26, 2029	DP			
	10517507	Jun 13, 2032	DP			
	11229378	Jul 11, 2031	DP			
	11464423	Sep 15, 2030	DP			
	11476952	Apr 28, 2026	DP			
	7978064	Sep 14, 2026	DP			
	8114021	Jun 21, 2030	DP			
	8258962	Nov 25, 2030	DP			
	8545402	Apr 27, 2030	DP			
	8547248	Dec 18, 2030	DP U-2167			
	8674825	Apr 09, 2029	DP U-2170			
	8718193	Dec 05, 2029	DP			
	8759350	Mar 02, 2027	U-1529			
	8847766	Mar 29, 2030	DP U-2167			
	8945005	Aug 19, 2029	DP U-2167			
	8956288	Jul 06, 2029	DP U-2167			
	8961412	Nov 17, 2030	DP			
	9060708	Mar 05, 2029	DP			
	9119554	Dec 16, 2028	DP			
	9125939	Jul 28, 2026	U-1749			
	9149577	Dec 15, 2029	DP			
	9258035	Mar 05, 2029	DP			
	9268909	Oct 15, 2033	DP U-2168			
	9320455	Dec 15, 2031	DP			
	9433371	Sep 15, 2029	DP			
	9444503	Nov 19, 2027	DP U-2169			
	9941931	Nov 04, 2030	DP			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>ARIPIRAZOLE - ABILIFY MYCITE KIT</u>						
N 207202 003	10441194	Jul 26, 2029	DP			
	10517507	Jun 13, 2032	DP			
	11229378	Jul 11, 2031	DP			
	11464423	Sep 15, 2030	DP			
	11476952	Apr 28, 2026	DP			
	7978064	Sep 14, 2026	DP			
	8114021	Jun 21, 2030	DP			
	8258962	Nov 25, 2030	DP			
	8545402	Apr 27, 2030	DP			
	8547248	Dec 18, 2030	DP	U-2167		
	8674825	Apr 09, 2029	DP	U-2170		
	8718193	Dec 05, 2029	DP			
	8759350	Mar 02, 2027		U-1529		
	8847766	Mar 29, 2030	DP	U-2167		
	8945005	Aug 19, 2029	DP	U-2167		
	8956288	Jul 06, 2029	DP	U-2167		
	8961412	Nov 17, 2030	DP			
	9060708	Mar 05, 2029	DP			
	9119554	Dec 16, 2028	DP			
	9125939	Jul 28, 2026		U-1749		
	9149577	Dec 15, 2029	DP			
	9258035	Mar 05, 2029	DP			
	9268909	Oct 15, 2033	DP	U-2168		
	9320455	Dec 15, 2031	DP			
	9433371	Sep 15, 2029	DP			
	9444503	Nov 19, 2027	DP	U-2169		
	9941931	Nov 04, 2030	DP			
<u>ARIPIRAZOLE - ABILIFY MYCITE KIT</u>						
N 207202 004	10441194	Jul 26, 2029	DP			
	10517507	Jun 13, 2032	DP			
	11229378	Jul 11, 2031	DP			
	11464423	Sep 15, 2030	DP			
	11476952	Apr 28, 2026	DP			
	7978064	Sep 14, 2026	DP			
	8114021	Jun 21, 2030	DP			
	8258962	Nov 25, 2030	DP			
	8545402	Apr 27, 2030	DP			
	8547248	Dec 18, 2030	DP	U-2167		
	8674825	Apr 09, 2029	DP	U-2170		
	8718193	Dec 05, 2029	DP			
	8759350	Mar 02, 2027		U-1529		
	8847766	Mar 29, 2030	DP	U-2167		
	8945005	Aug 19, 2029	DP	U-2167		
	8956288	Jul 06, 2029	DP	U-2167		
	8961412	Nov 17, 2030	DP			
	9060708	Mar 05, 2029	DP			
	9119554	Dec 16, 2028	DP			
	9125939	Jul 28, 2026		U-1749		
	9149577	Dec 15, 2029	DP			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>ARIPIRAZOLE - ABILIFY MYCITE KIT</u>						
N 207202 004	9258035	Mar 05, 2029	DP			
	9268909	Oct 15, 2033	DP U-2168			
	9320455	Dec 15, 2031	DP			
	9433371	Sep 15, 2029	DP			
	9444503	Nov 19, 2027	DP U-2169			
	9941931	Nov 04, 2030	DP			
<u>ARIPIRAZOLE - ABILIFY MYCITE KIT</u>						
N 207202 005	10441194	Jul 26, 2029	DP			
	10517507	Jun 13, 2032	DP			
	11229378	Jul 11, 2031	DP			
	11464423	Sep 15, 2030	DP			
	11476952	Apr 28, 2026	DP			
	7978064	Sep 14, 2026	DP			
	8114021	Jun 21, 2030	DP			
	8258962	Nov 25, 2030	DP			
	8545402	Apr 27, 2030	DP			
	8547248	Dec 18, 2030	DP U-2167			
	8674825	Apr 09, 2029	DP U-2170			
	8718193	Dec 05, 2029	DP			
	8759350	Mar 02, 2027	U-1529			
	8847766	Mar 29, 2030	DP U-2167			
	8945005	Aug 19, 2029	DP U-2167			
	8956288	Jul 06, 2029	DP U-2167			
	8961412	Nov 17, 2030	DP			
	9060708	Mar 05, 2029	DP			
	9119554	Dec 16, 2028	DP			
	9125939	Jul 28, 2026	U-1749			
	9149577	Dec 15, 2029	DP			
	9258035	Mar 05, 2029	DP			
	9268909	Oct 15, 2033	DP U-2168			
	9320455	Dec 15, 2031	DP			
	9433371	Sep 15, 2029	DP			
	9444503	Nov 19, 2027	DP U-2169			
	9941931	Nov 04, 2030	DP			
<u>ARIPIRAZOLE - ABILIFY MYCITE KIT</u>						
N 207202 006	10441194	Jul 26, 2029	DP			
	10517507	Jun 13, 2032	DP			
	11229378	Jul 11, 2031	DP			
	11464423	Sep 15, 2030	DP			
	11476952	Apr 28, 2026	DP			
	7978064	Sep 14, 2026	DP			
	8114021	Jun 21, 2030	DP			
	8258962	Nov 25, 2030	DP			
	8545402	Apr 27, 2030	DP			
	8547248	Dec 18, 2030	DP U-2167			
	8674825	Apr 09, 2029	DP U-2170			
	8718193	Dec 05, 2029	DP			
	8759350	Mar 02, 2027	U-1529			
	8847766	Mar 29, 2030	DP U-2167			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>ARIPIRAZOLE - ABILIFY MYCITE KIT</u>						
N 207202	006	8945005	Aug 19, 2029	DP	U-2167	
		8956288	Jul 06, 2029	DP	U-2167	
		8961412	Nov 17, 2030	DP		
		9060708	Mar 05, 2029	DP		
		9119554	Dec 16, 2028	DP		
		9125939	Jul 28, 2026		U-1749	
		9149577	Dec 15, 2029	DP		
		9258035	Mar 05, 2029	DP		
		9268909	Oct 15, 2033	DP	U-2168	
		9320455	Dec 15, 2031	DP		
		9433371	Sep 15, 2029	DP		
		9444503	Nov 19, 2027	DP	U-2169	
		9941931	Nov 04, 2030	DP		
<u>ARIPIRAZOLE - OPIPZA</u>						
N 216655	001	11331315	Nov 03, 2040	DP		
		11701352	Dec 15, 2041	DP		
<u>ARIPIRAZOLE - OPIPZA</u>						
N 216655	002	11331315	Nov 03, 2040	DP		
		11701352	Dec 15, 2041	DP		
<u>ARIPIRAZOLE - OPIPZA</u>						
N 216655	003	11331315	Nov 03, 2040	DP		
		11701352	Dec 15, 2041	DP		
<u>ARIPIRAZOLE - ABILIFY ASIMTUFII</u>						
N 217006	001	10517951	Apr 23, 2033	DP	U-3245	
		10517951	Apr 23, 2033	DP	U-814	
		11097007	Apr 23, 2033	DP	U-3245	
		11097007	Apr 23, 2033	DP	U-814	
		11638757	Apr 23, 2033	DP	U-3245	
		11638757	Apr 23, 2033	DP	U-814	
		12016927	Apr 23, 2033	DP	U-3245	
		12016927	Apr 23, 2033	DP	U-543	
		8338427	Mar 15, 2025	DP	U-1530	
		8399469	Jun 29, 2025	DS		
<u>ARIPIRAZOLE - ABILIFY ASIMTUFII</u>						
N 217006	002	10517951	Apr 23, 2033	DP	U-3245	
		10517951	Apr 23, 2033	DP	U-814	
		11097007	Apr 23, 2033	DP	U-3245	
		11097007	Apr 23, 2033	DP	U-814	
		11638757	Apr 23, 2033	DP	U-3245	
		11638757	Apr 23, 2033	DP	U-814	
		12016927	Apr 23, 2033	DP	U-3245	
		12016927	Apr 23, 2033	DP	U-543	
		8338427	Mar 15, 2025	DP	U-1530	
		8399469	Jun 29, 2025	DS		
<u>ARIPIRAZOLE LAUROXIL - ARISTADA</u>						
N 207533	001	10112903	Jun 24, 2030	DS	U-543	
		10226458	Mar 19, 2032		U-543	

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>ARIPIRAZOLE LAUROXIL - ARISTADA</u>						
N 207533 001	10238651	Mar 19, 2035	U-2402			
	10813928	Mar 19, 2035	U-2983			
	11097006	Oct 24, 2033	DP U-764			
	11273158	Apr 06, 2039	U-543			
	11406632	Mar 19, 2035	U-2402			
	8431576	Oct 26, 2030	DS			
	8796276	Jun 24, 2030	U-543			
	9034867	Nov 07, 2032	DP U-543			
	9193685	Oct 24, 2033	DP U-543			
	9452131	Mar 19, 2035	U-2402			
<u>ARIPIRAZOLE LAUROXIL - ARISTADA</u>						
N 207533 002	10112903	Jun 24, 2030	DS U-543			
	10226458	Mar 19, 2032	U-543			
	10238651	Mar 19, 2035	U-2402			
	10813928	Mar 19, 2035	U-2983			
	11097006	Oct 24, 2033	DP U-764			
	11273158	Apr 06, 2039	U-543			
	11406632	Mar 19, 2035	U-2402			
	8431576	Oct 26, 2030	DS			
	8796276	Jun 24, 2030	U-543			
	9034867	Nov 07, 2032	DP U-543			
	9193685	Oct 24, 2033	DP U-543			
	9452131	Mar 19, 2035	U-2402			
	9526726	Mar 19, 2035	DP			
<u>ARIPIRAZOLE LAUROXIL - ARISTADA</u>						
N 207533 003	10112903	Jun 24, 2030	DS U-543			
	10226458	Mar 19, 2032	U-543			
	10238651	Mar 19, 2035	U-2402			
	10813928	Mar 19, 2035	U-2402			
	11097006	Oct 24, 2033	DP U-764			
	11273158	Apr 06, 2039	U-543			
	11406632	Mar 19, 2035	U-2402			
	8431576	Oct 26, 2030	DS			
	8796276	Jun 24, 2030	U-543			
	9034867	Nov 07, 2032	DP U-543			
	9193685	Oct 24, 2033	DP U-543			
	9452131	Mar 19, 2035	U-2402			
	9526726	Mar 19, 2035	DP			
<u>ARIPIRAZOLE LAUROXIL - ARISTADA</u>						
N 207533 004	10112903	Jun 24, 2030	DS U-543			
	10226458	Mar 19, 2032	U-543			
	10238651	Mar 19, 2035	U-2402			
	10813928	Mar 19, 2035	U-2983			
	11097006	Oct 24, 2033	DP U-764			
	11273158	Apr 06, 2039	U-543			
	11406632	Mar 19, 2035	U-2402			
	8431576	Oct 26, 2030	DS			
	8796276	Jun 24, 2030	U-543			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>ARIPIRAZOLE LAUROXIL - ARISTADA</u>						
N 207533	004 9034867	Nov 07, 2032	DP	U-543		
	9193685	Oct 24, 2033	DP	U-543		
	9452131	Mar 19, 2035		U-2402		
<u>ARIPIRAZOLE LAUROXIL - ARISTADA INITIO KIT</u>						
N 209830	001 10016415	Sep 08, 2035	DP			
	10112903	Jun 24, 2030	DS	U-543		
	10688091	Aug 17, 2035	DP			
	10849894	Aug 17, 2035		U-543		
	11154552	Aug 17, 2035	DP			
	11273158	Apr 06, 2039		U-543		
	8431576	Oct 26, 2030	DS			
	8796276	Jun 24, 2030		U-543		
<u>ARSENIC TRIOXIDE - TRISENOX</u>						
N 021248	001				ODE-167	Jan 12, 2025
<u>ARSENIC TRIOXIDE - TRISENOX</u>						
N 021248	002				ODE-167	Jan 12, 2025
<u>ARTESUNATE - ARTESUNATE</u>						
N 213036	001 12121506	Feb 16, 2044	DP		NCE	May 26, 2025
					ODE-290	May 26, 2027
<u>ASCIMINIB HYDROCHLORIDE - SCEMBLIX</u>						
N 215358	001 11407735	May 14, 2040	DS		I-953	Oct 29, 2027
	8829195	May 13, 2033	DS	U-1374	I-954	Oct 29, 2027
					NCE	Oct 29, 2026
					ODE-381	Oct 29, 2028
					ODE-382	Oct 29, 2028
					ODE-499	Oct 29, 2031
					ODE-500	Oct 29, 2031
<u>ASCIMINIB HYDROCHLORIDE - SCEMBLIX</u>						
N 215358	002 11407735	May 14, 2040	DS		I-953	Oct 29, 2027
	8829195	May 13, 2033	DS	U-1374	I-954	Oct 29, 2027
					NCE	Oct 29, 2026
					ODE-381	Oct 29, 2028
					ODE-382	Oct 29, 2028
					ODE-499	Oct 29, 2031
					ODE-500	Oct 29, 2031
<u>ASCIMINIB HYDROCHLORIDE - SCEMBLIX</u>						
N 215358	003 11407735	May 14, 2040	DS		NCE	Oct 29, 2026
	8829195	May 13, 2033	DS	U-1374	ODE*	Oct 29, 2028
<u>ASCORBIC ACID; POLYETHYLENE GLYCOL 3350; POTASSIUM CHLORIDE; SODIUM ASCORBATE; SODIUM CHLORIDE; SODIUM SULFATE - PLENVU</u>						
N 209381	001 10016504	Sep 10, 2033	DP			
	10646512	Mar 25, 2032	DP			
	10780112	Mar 09, 2032	DP			
	10792306	Mar 09, 2032	DP	U-2310		
	10918723	Sep 10, 2033		U-2310		
	11529368	Mar 09, 2032	DP	U-2310		
	12083179	Sep 10, 2033	DP	U-2310		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>ASCORBIC ACID; POLYETHYLENE GLYCOL 3350; POTASSIUM CHLORIDE; SODIUM ASCORBATE; SODIUM CHLORIDE; SODIUM SULFATE - PLENVU</u>						
N 209381	001	8999313	Sep 10, 2033	DP		
		9326969	Sep 10, 2033		U-2310	
		9592252	Aug 11, 2032	DP	U-2310	
		9707297	Sep 10, 2033	DP		
<u>ASENAPINE - SECUADO</u>						
N 212268	001	10022445	Jul 25, 2033	DP		
		10583121	Jul 25, 2033	DP	U-2763	
		10814002	Jul 25, 2033	DP	U-2763	
		11123305	Jul 25, 2033	DP		
		11813364	Sep 22, 2033	DP		
		9687474	Jul 25, 2033	DP		
<u>ASENAPINE - SECUADO</u>						
N 212268	002	10022445	Jul 25, 2033	DP		
		10583121	Jul 25, 2033	DP	U-2763	
		10814002	Jul 25, 2033	DP	U-2763	
		11123305	Jul 25, 2033	DP		
		11813364	Sep 22, 2033	DP		
		9687474	Jul 25, 2033	DP		
<u>ASENAPINE - SECUADO</u>						
N 212268	003	10022445	Jul 25, 2033	DP		
		10583121	Jul 25, 2033	DP	U-2763	
		10814002	Jul 25, 2033	DP	U-2763	
		11123305	Jul 25, 2033	DP		
		11813364	Sep 22, 2033	DP		
		9687474	Jul 25, 2033	DP		
<u>ASENAPINE MALEATE - SAPHRIS</u>						
N 022117	001	7741358	Apr 06, 2026	DS DP	U-1064	
		7741358	Apr 06, 2026	DS DP	U-1960	
		7741358	Apr 06, 2026	DS DP	U-1961	
		7741358	Apr 06, 2026	DS DP	U-1962	
		7741358	Apr 06, 2026	DS DP	U-1963	
		7741358*PED	Oct 06, 2026			
		8022228	Apr 06, 2026	DS DP		
		8022228*PED	Oct 06, 2026			
<u>ASENAPINE MALEATE - SAPHRIS</u>						
N 022117	002	7741358	Apr 06, 2026	DS DP	U-1064	
		7741358	Apr 06, 2026	DS DP	U-1960	
		7741358	Apr 06, 2026	DS DP	U-1961	
		7741358	Apr 06, 2026	DS DP	U-1962	
		7741358	Apr 06, 2026	DS DP	U-1963	
		7741358*PED	Oct 06, 2026			
		8022228	Apr 06, 2026	DS DP		
		8022228*PED	Oct 06, 2026			
<u>ASENAPINE MALEATE - SAPHRIS</u>						
N 022117	003	7741358	Apr 06, 2026	DS DP	U-1893	
		7741358	Apr 06, 2026	DS DP	U-1966	
		7741358*PED	Oct 06, 2026			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>ASENAPINE MALEATE - SAPHRIS</u>						
N 022117	003 8022228	Apr 06, 2026	DS DP			
	8022228*PED	Oct 06, 2026				
<u>ASPIRIN - VAZALORE</u>						
N 203697	001 10646431	Sep 29, 2032	DP			
	10786444	Sep 29, 2032		U-3559		
	9216150	Sep 29, 2032	DP			
	9226892	Sep 29, 2032		U-1731		
	9226892	Sep 29, 2032		U-1732		
	9226892	Sep 29, 2032		U-1733		
<u>ASPIRIN - VAZALORE</u>						
N 203697	002 10646431	Sep 29, 2032	DP			
	10786444	Sep 29, 2032		U-3559		
<u>ASPIRIN; OMEPRAZOLE - YOSPRALA</u>						
N 205103	001 9539214	Mar 13, 2033		U-1902		
	9987231	Jan 02, 2033		U-2324		
<u>ASPIRIN; OMEPRAZOLE - YOSPRALA</u>						
N 205103	002 9539214	Mar 13, 2033		U-1902		
	9987231	Jan 02, 2033		U-2324		
<u>ATAZANAVIR SULFATE; COBICISTAT - EVOTAZ</u>						
N 206353	001 10039718	Oct 06, 2032	DP			
	8148374	Sep 03, 2029	DS DP	U-1279		
<u>ATOGEFANT - OULIPTA</u>						
N 215206	001 10117836	Jan 30, 2035	DP		I-909	Apr 17, 2026
	8754096	Jul 19, 2032	DS DP	U-3534	NCE	Sep 28, 2026
	9499545	Nov 10, 2031	DS DP	U-3534		
	9850246	Mar 13, 2033	DS			
<u>ATOGEFANT - OULIPTA</u>						
N 215206	002 10117836	Jan 30, 2035	DP		I-909	Apr 17, 2026
	8754096	Jul 19, 2032	DS DP	U-3534	NCE	Sep 28, 2026
	9499545	Nov 10, 2031	DS DP	U-3534		
	9850246	Mar 13, 2033	DS			
<u>ATOGEFANT - OULIPTA</u>						
N 215206	003 10117836	Jan 30, 2035	DP		I-909	Apr 17, 2026
	12090148	Jul 29, 2041		U-3534	NCE	Sep 28, 2026
	8754096	Jul 19, 2032	DS DP	U-3534		
	9499545	Nov 10, 2031	DS DP	U-3534		
	9850246	Mar 13, 2033	DS			
<u>ATORVASTATIN CALCIUM - ATORVALIO</u>						
N 213260	001 11369567	Jun 07, 2037	DP			
	11654106	Jun 07, 2037	DP	U-3612		
	11654106	Jun 07, 2037	DP	U-3613		
	11925704	Jun 07, 2037	DP	U-3853		
	12168069	Jun 07, 2037	DP			
<u>AVACINCAPTAD PEGOL SODIUM - IZERVAY</u>						
N 217225	001 10947544	Feb 14, 2025	DS	U-3673		
	11273171	Jul 11, 2034		U-3673		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>AVACINCAPTAD PEGOL SODIUM - IZERVAY</u>						
N 217225	001	11491176	Jul 11, 2034	U-3673		
		12016875	Jul 11, 2034	U-3673		
		7538211	Feb 14, 2025	DS		
		7579456	Feb 14, 2025	DS		
		7803931	Feb 14, 2025	DS		
		8236773	Nov 11, 2026	U-3673		
		9617546	Feb 14, 2025	DS	U-3673	
<u>AVACOPAN - TAVNEOS</u>						
N 214487	001	11603356	May 29, 2041	DS DP U-3558	NCE	Oct 07, 2026
		11951214	Nov 27, 2039	DP	ODE-377	Oct 07, 2028
		8445515	Feb 03, 2031	DS DP		
		8906938	Dec 21, 2029	DS DP		
<u>AVANAFIL - STENDRA</u>						
N 202276	001	6656935	Apr 27, 2025	DS DP U-155	M-282	Oct 18, 2025
<u>AVANAFIL - STENDRA</u>						
N 202276	002	6656935	Apr 27, 2025	DS DP U-155	M-282	Oct 18, 2025
<u>AVANAFIL - STENDRA</u>						
N 202276	003	6656935	Apr 27, 2025	DS DP U-155	M-282	Oct 18, 2025
<u>AVAPRITINIB - AYPAKIT</u>						
N 212608	001	11964980	Apr 10, 2040	DS DP	NCE	Jan 09, 2025
		11999744	Apr 10, 2040	U-3168	ODE-356	Jun 16, 2028
		12060354	Mar 08, 2042	U-2726	ODE-366	Jan 09, 2027
		9200002	Oct 15, 2034	DS DP U-2726	ODE-434	May 22, 2030
		9200002	Oct 15, 2034	DS DP U-3168		
		9944651	Oct 15, 2034	DS DP U-2726		
		9944651	Oct 15, 2034	DS DP U-3168		
		9994575	Oct 15, 2034	DS DP U-2726		
		9994575	Oct 15, 2034	DS DP U-3168		
<u>AVAPRITINIB - AYPAKIT</u>						
N 212608	002	11964980	Apr 10, 2040	DS DP	NCE	Jan 09, 2025
		11999744	Apr 10, 2040	U-3168	ODE-356	Jun 16, 2028
		12060354	Mar 08, 2042	U-2726	ODE-366	Jan 09, 2027
		9200002	Oct 15, 2034	DS DP U-2726	ODE-434	May 22, 2030
		9200002	Oct 15, 2034	DS DP U-3168		
		9944651	Oct 15, 2034	DS DP U-2726		
		9944651	Oct 15, 2034	DS DP U-3168		
		9994575	Oct 15, 2034	DS DP U-2726		
		9994575	Oct 15, 2034	DS DP U-3168		
<u>AVAPRITINIB - AYPAKIT</u>						
N 212608	003	11964980	Apr 10, 2040	DS DP	NCE	Jan 09, 2025
		12060354	Mar 08, 2042	U-2726	ODE-356	Jun 16, 2028
		9200002	Oct 15, 2034	DS DP U-2726	ODE-366	Jan 09, 2027
		9944651	Oct 15, 2034	DS DP U-2726	ODE-434	May 22, 2030
		9994575	Oct 15, 2034	DS DP U-2726		
<u>AVAPRITINIB - AYPAKIT</u>						
N 212608	004	11827642	Oct 15, 2034	DS DP U-3506	I-912	May 22, 2026

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>AVAPRITINIB - AYVAKIT</u>						
N 212608 004	11964980	Apr 10, 2040	DS DP		NCE	Jan 09, 2025
	11999744	Apr 10, 2040	U-3168		ODE-356	Jun 16, 2028
	11999744	Apr 10, 2040	U-3506		ODE-366	Jan 09, 2027
	9200002	Oct 15, 2034	DS DP U-3168		ODE-434	May 22, 2030
	9200002	Oct 15, 2034	DS DP U-3506			
	9944651	Oct 15, 2034	DS DP U-3168			
	9944651	Oct 15, 2034	DS DP U-3506			
	9994575	Oct 15, 2034	DS DP U-3168			
	9994575	Oct 15, 2034	DS DP U-3506			
<u>AVAPRITINIB - AYVAKIT</u>						
N 212608 005	11964980	Apr 10, 2040	DS DP		NCE	Jan 09, 2025
	11999744	Apr 10, 2040	U-3168		ODE-356	Jun 16, 2028
	9200002	Oct 15, 2034	DS DP U-3168		ODE-366	Jan 09, 2027
	9944651	Oct 15, 2034	DS DP U-3168		ODE-434	May 22, 2030
	9994575	Oct 15, 2034	DS DP U-3168			
<u>AVATROMBOPAG MALEATE - DOPTOLET</u>						
N 210238 001	7638536	Jul 28, 2027	DS DP		ODE-246	Jun 26, 2026
<u>AVIBACTAM SODIUM; CEFTAZIDIME - AVYCAZ</u>						
N 206494 001	7112592	Jan 07, 2026	DS DP U-2244		NCE	Feb 25, 2020
	7112592	Jan 07, 2026	DS DP U-2508		NPP	Dec 20, 2025
	7112592	Jan 07, 2026	DS DP U-282		NPP	Jan 26, 2027
	7112592	Jan 07, 2026	DS DP U-3818		GAIN	Feb 25, 2025
	7612087	Nov 12, 2026	DP			
	8471025	Aug 12, 2031	DS			
	8835455	Oct 08, 2030	DP			
	8969566	Jun 15, 2032	DS			
	9284314	Jun 15, 2032	DS			
	9695122	Jun 15, 2032	DS			
<u>AXITINIB - INLYTA</u>						
N 202324 001	10570202	Feb 03, 2035	U-2844			
	10570202*PED	Aug 03, 2035				
	10869924	Jan 12, 2037	U-3044			
	10869924*PED	Jul 12, 2037				
	6534524	Apr 29, 2025	DS DP			
	6534524*PED	Oct 29, 2025				
	8791140	Dec 14, 2030	DS			
	8791140*PED	Jun 14, 2031				
<u>AXITINIB - INLYTA</u>						
N 202324 002	10570202	Feb 03, 2035	U-2844			
	10570202*PED	Aug 03, 2035				
	10869924	Jan 12, 2037	U-3044			
	10869924*PED	Jul 12, 2037				
	6534524	Apr 29, 2025	DS DP			
	6534524*PED	Oct 29, 2025				
	8791140	Dec 14, 2030	DS			
	8791140*PED	Jun 14, 2031				

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>AZACITIDINE - VIDAZA</u>						
N 050794	001				I-889	May 20, 2025
					ODE-399	May 20, 2029
<u>AZACITIDINE - ONUREG</u>						
N 214120	001	11571436	May 14, 2029	DP	ODE-320	Sep 01, 2027
		12053482	May 14, 2029	U-2950		
		8846628	Jun 03, 2030	DP U-2950		
<u>AZACITIDINE - ONUREG</u>						
N 214120	002	11571436	May 14, 2029	DP	ODE-320	Sep 01, 2027
		12053482	May 14, 2029	U-2950		
		8846628	Jun 03, 2030	DP U-2950		
<u>AZELAIC ACID - FINACEA</u>						
N 207071	001	10117812	Oct 18, 2027	DP U-1796		
		7700076	Sep 18, 2027	DP		
		9211259	Feb 28, 2029	U-1796		
		9265725	Dec 08, 2027	DP		
<u>AZELASTINE HYDROCHLORIDE - ASTEPRO</u>						
N 022203	001	8071073	Jun 04, 2028	DP		
		8518919	Nov 22, 2025	U-1430		
<u>AZELASTINE HYDROCHLORIDE - ASTEPRO</u>						
N 022203	002	8071073	Jun 04, 2028	DP		
		8518919	Nov 22, 2025	U-1430		
		9919050	Nov 22, 2025	DP		
<u>AZELASTINE HYDROCHLORIDE - ASTEPRO ALLERGY</u>						
N 213872	001	8071073	Jun 04, 2028	DP		
		8518919	Nov 22, 2025	U-3166		
		9919050	Nov 22, 2025	DP		
<u>AZELASTINE HYDROCHLORIDE - CHILDREN'S ASTEPRO ALLERGY</u>						
N 213872	002	8071073	Jun 04, 2028	DP		
		8518919	Nov 22, 2025	U-3166		
		9919050	Nov 22, 2025	DP		
<u>AZELASTINE HYDROCHLORIDE; FLUTICASONE PROPIONATE - DYMISTA</u>						
N 202236	001	8168620	Feb 24, 2026	DP		
<u>AZILSARTAN KAMEDOXOMIL - EDARBI</u>						
N 200796	001	7157584	May 22, 2025	DS		
		7572920	Jan 07, 2025	DP U-3		
		9066936	Mar 26, 2028	DP		
<u>AZILSARTAN KAMEDOXOMIL - EDARBI</u>						
N 200796	002	7157584	May 22, 2025	DS		
		7572920	Jan 07, 2025	DP U-3		
		9066936	Mar 26, 2028	DP		
<u>AZILSARTAN KAMEDOXOMIL; CHLORTHALIDONE - EDARBYCLOR</u>						
N 202331	001	7157584	May 22, 2025	DS		
		7572920	Jan 07, 2025	DP U-3		
		9066936	Mar 26, 2028	DP		
		9169238	Feb 04, 2030	DP		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>AZILSARTAN KAMEDOXOMIL; CHLORTHALIDONE - EDARBYCLOR</u>						
N 202331	001 9387249	Jul 01, 2031	U-3			
<u>AZILSARTAN KAMEDOXOMIL; CHLORTHALIDONE - EDARBYCLOR</u>						
N 202331	002 7157584	May 22, 2025	DS			
	7572920	Jan 07, 2025	DP U-3			
	9066936	Mar 26, 2028	DP			
	9169238	Feb 04, 2030	DP			
	9387249	Jul 01, 2031	U-3			
<u>BACLOFEN - OZOBAX</u>						
N 208193	001 10610502	Aug 30, 2039	U-2779			
<u>BACLOFEN - LYVISPAH</u>						
N 215422	001 10792262	Jul 29, 2039	DP U-3263			
	11491125	Sep 29, 2041	DP U-3488			
	11491125	Sep 29, 2041	DP U-3489			
	11654124	Jul 29, 2039	DP			
	11850225	Sep 29, 2041	DP U-3488			
	11850225	Sep 29, 2041	DP U-3489			
	11931328	Jul 29, 2039	DP			
<u>BACLOFEN - LYVISPAH</u>						
N 215422	002 10792262	Jul 29, 2039	DP U-3263			
	11491125	Sep 29, 2041	DP U-3488			
	11491125	Sep 29, 2041	DP U-3489			
	11654124	Jul 29, 2039	DP			
	11850225	Sep 29, 2041	DP U-3488			
	11850225	Sep 29, 2041	DP U-3489			
	11931328	Jul 29, 2039	DP			
<u>BACLOFEN - LYVISPAH</u>						
N 215422	003 10792262	Jul 29, 2039	DP U-3263			
	11491125	Sep 29, 2041	DP U-3488			
	11491125	Sep 29, 2041	DP U-3489			
	11654124	Jul 29, 2039	DP			
	11850225	Sep 29, 2041	DP U-3488			
	11850225	Sep 29, 2041	DP U-3489			
	11931328	Jul 29, 2039	DP			
<u>BACLOFEN - FLEOSUVY</u>						
N 215602	001 11324696	Sep 29, 2037	DP			
	11446246	Sep 08, 2037	U-3433			
<u>BALOXAVIR MARBOXIL - XOFLUZA</u>						
N 210854	001 10392406	Apr 27, 2036	DS			
	10633397	Apr 27, 2036	U-2816			
	10633397	Apr 27, 2036	U-3000			
	10759814	Aug 09, 2037	DS DP			
	11261198	Sep 25, 2038	DP			
	11306106	Aug 09, 2037	U-2816			
	11306106	Aug 09, 2037	U-3000			
	12064438	Oct 09, 2039	DP			
	8927710	May 05, 2031	DP			
	8987441	Sep 21, 2031	DS DP			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>BALOXAVIR MARBOXIL - XOFLUZA</u>						
N 210854	001	9815835	Jun 14, 2030	DP		
<u>BALOXAVIR MARBOXIL - XOFLUZA</u>						
N 210854	002	10392406	Apr 27, 2036	DS		
		10633397	Apr 27, 2036		U-2816	
		10633397	Apr 27, 2036		U-3000	
		10759814	Aug 09, 2037	DS DP		
		11261198	Sep 25, 2038	DP		
		11306106	Aug 09, 2037		U-2816	
		11306106	Aug 09, 2037		U-3000	
		12064438	Oct 09, 2039	DP		
		8927710	May 05, 2031	DP		
		8987441	Sep 21, 2031	DS DP		
		9815835	Jun 14, 2030	DP		
<u>BALOXAVIR MARBOXIL - XOFLUZA</u>						
N 210854	003	10392406	Apr 27, 2036	DS		
		10633397	Apr 27, 2036		U-2816	
		10633397	Apr 27, 2036		U-3000	
		10759814	Aug 09, 2037	DS DP		
		11261198	Sep 25, 2038	DP		
		11306106	Aug 09, 2037		U-2816	
		11306106	Aug 09, 2037		U-3000	
		12064438	Oct 09, 2039	DP		
		8927710	May 05, 2031	DP		
		8987441	Sep 21, 2031	DS DP		
		9815835	Jun 14, 2030	DP		
<u>BALOXAVIR MARBOXIL - XOFLUZA</u>						
N 214410	001	10392406	Apr 27, 2036	DS		
		10633397	Apr 27, 2036		U-2816	
		10633397	Apr 27, 2036		U-3000	
		10759814	Aug 09, 2037	DS DP		
		11261198	Sep 25, 2038	DP		
		11306106	Aug 09, 2037		U-2816	
		11306106	Aug 09, 2037		U-3000	
		11925648	Apr 21, 2041	DP		
		8927710	May 05, 2031	DP		
		8987441	Sep 21, 2031	DS DP		
		9815835	Jun 14, 2030	DP		
<u>BALSALAZIDE DISODIUM - COLAZAL</u>						
N 020610	001	7452872	Aug 24, 2026		U-141	
		7625884	Aug 24, 2026		U-141	
<u>BALSALAZIDE DISODIUM - GIAZO</u>						
N 022205	001	7452872	Aug 24, 2026		U-1229	
		7625884	Aug 24, 2026		U-1229	
		8497256	Jun 23, 2031		U-1229	
		9192616	Aug 02, 2026		U-1229	

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>BARICITINIB - OLUMIANT</u>						
N 207924 001	11045474	Nov 30, 2032	U-3372		I-890	Jun 13, 2025
	11806555	Nov 02, 2031	U-3500		I-891	May 10, 2025
	8158616	May 31, 2032	DS DP			
	8420629	Mar 10, 2029	U-247			
	9089574	Nov 30, 2032	U-3372			
	9737469	Nov 02, 2031	U-3500			
<u>BARICITINIB - OLUMIANT</u>						
N 207924 002	11045474	Nov 30, 2032	U-3372		I-890	Jun 13, 2025
	11806555	Nov 02, 2031	U-3500		I-891	May 10, 2025
	8158616	May 31, 2032	DS DP			
	8420629	Mar 10, 2029	U-247			
	9089574	Nov 30, 2032	U-3372			
	9737469	Nov 02, 2031	U-3500			
<u>BARICITINIB - OLUMIANT</u>						
N 207924 003	11045474	Nov 30, 2032	U-3372		I-890	Jun 13, 2025
	11806555	Nov 02, 2031	U-3500		I-891	May 10, 2025
	8158616	May 31, 2032	DS DP			
	9089574	Nov 30, 2032	U-3372			
	9737469	Nov 02, 2031	U-3500			
<u>BAZEDOXIFENE ACETATE; ESTROGENS, CONJUGATED - DUAVEE</u>						
N 022247 001	7683051	Mar 10, 2027	DS DP U-594			
	7683051	Mar 10, 2027	DS DP U-904			
<u>BECLOMETHASONE DIPROPIONATE - QVAR 80</u>						
N 020911 001	10022509	May 18, 2031	DP			
	10022510	May 18, 2031	DP			
	10086156	May 18, 2031	DP			
	10561808	Jan 01, 2032	DP			
	10695512	May 18, 2031	DP			
	11395889	May 18, 2031	DP			
	9463289	May 18, 2031	DP			
	9808587	May 18, 2031	DP			
<u>BECLOMETHASONE DIPROPIONATE - QVAR 40</u>						
N 020911 002	10022509	May 18, 2031	DP			
	10022510	May 18, 2031	DP			
	10086156	May 18, 2031	DP			
	10561808	Jan 01, 2032	DP			
	10695512	May 18, 2031	DP			
	11395889	May 18, 2031	DP			
	9463289	May 18, 2031	DP			
	9808587	May 18, 2031	DP			
<u>BECLOMETHASONE DIPROPIONATE - QNASL</u>						
N 202813 001	10188811	Oct 21, 2031	DP			
	7780038	Jan 24, 2027	DP			
<u>BECLOMETHASONE DIPROPIONATE - QNASL</u>						
N 202813 002	10188811	Oct 21, 2031	DP			
	7780038	Jan 24, 2027	DP			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>BECLOMETHASONE DIPROPIONATE - QVAR REDHALER</u>						
N 207921 001	10022509	May 18, 2031	DP			
	10022510	May 18, 2031	DP			
	10086156	May 18, 2031	DP			
	10561808	Jan 01, 2032	DP			
	10695512	May 18, 2031	DP			
	10792447	Jan 25, 2039	DP			
	11395888	Jan 26, 2038	DP			
	11395889	May 18, 2031	DP			
	11559637	Jul 21, 2039	DP			
	11583643	Aug 19, 2041	DP			
	11793953	Jan 26, 2038	DP			
	11865247	Jan 26, 2038	DP			
	11896759	Jan 26, 2038	DP			
	11957832	May 05, 2041	DP			
	8132712	Sep 07, 2028	DP			
	8931476	Jul 17, 2031	DP			
<u>BECLOMETHASONE DIPROPIONATE - QVAR REDHALER</u>						
N 207921 002	10022509	May 18, 2031	DP			
	10022510	May 18, 2031	DP			
	10086156	May 18, 2031	DP			
	10561808	Jan 01, 2032	DP			
	10695512	May 18, 2031	DP			
	10792447	Jan 25, 2039	DP			
	11395888	Jan 26, 2038	DP			
	11395889	May 18, 2031	DP			
	11559637	Jul 21, 2039	DP			
	11583643	Aug 19, 2041	DP			
	11793953	Jan 26, 2038	DP			
	11865247	Jan 26, 2038	DP			
	11896759	Jan 26, 2038	DP			
	11957832	May 05, 2041	DP			
	8132712	Sep 07, 2028	DP			
	8931476	Jul 17, 2031	DP			
<u>BEDAQUILINE FUMARATE - SIRTURO</u>						
N 204384 001	7498343	Dec 01, 2026	DS DP U-1321		M-306	Jun 21, 2027
	8546428	Mar 19, 2029	DS DP U-1321		ODE-251	Aug 09, 2026
					ODE-307	May 27, 2027
<u>BEDAQUILINE FUMARATE - SIRTURO</u>						
N 204384 002	7498343	Dec 01, 2026	DS DP U-1321		M-306	Jun 21, 2027
	8546428	Mar 19, 2029	DS DP U-1321		ODE-307	May 27, 2027
<u>BELINOSTAT - BELEODAQ</u>						
N 206256 001	6888027	Aug 10, 2026	DS DP U-1544			
	8835501	Oct 27, 2027	DP			
<u>BELUMOSUDIL MESYLATE - REZUROCK</u>						
N 214783 001	10183931	Oct 07, 2033	U-3246		NCE	Jul 16, 2026
	10696660	Oct 07, 2033	U-3246		ODE-362	Jul 16, 2028
	11311541	Apr 09, 2035	U-3369			
	12097202	Jul 14, 2042	U-4014			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>BELUMOSUDIL MESYLATE - REZUROCK</u>						
N 214783	001 8357693	Oct 30, 2029	DS DP U-3247			
<u>BELZUTIFAN - WELIREG</u>						
N 215383	001 9908845	Sep 05, 2034	DS DP U-3201		I-931	Dec 14, 2026
	9908845	Sep 05, 2034	DS DP U-3780		NCE	Aug 13, 2026
	RE49948	Sep 05, 2034	DS DP U-3201		ODE-364	Aug 13, 2028
	RE49948	Sep 05, 2034	DS DP U-3780			
<u>BEMPEDOIC ACID - NEXLETOI</u>						
N 211616	001 11613511	Jun 19, 2040	DS		I-943	Mar 22, 2027
	11744816	Mar 14, 2036	U-3883		I-944	Mar 22, 2027
	11760714	Jun 19, 2040	DP		NCE	Feb 21, 2025
	11926584	Jun 19, 2040	U-3873			
	7335799	Dec 03, 2030	DS			
<u>BEMPEDOIC ACID; EZETIMIBE - NEXLIZET</u>						
N 211617	001 10912751	Mar 14, 2036	U-3224		I-943	Mar 22, 2027
	10912751	Mar 14, 2036	U-3884		I-945	Mar 22, 2027
	11613511	Jun 19, 2040	DS		NCE	Feb 21, 2025
	11744816	Mar 14, 2036	U-3692			
	11744816	Mar 14, 2036	U-3883			
	11760714	Jun 19, 2040	DP			
	11926584	Jun 19, 2040	U-3873			
	7335799	Dec 03, 2030	DS			
<u>BENDAMUSTINE HYDROCHLORIDE - TREANDA</u>						
N 022249	001 8436190	Oct 26, 2030	DP			
	8436190*PED	Apr 26, 2031				
	8445524	Mar 26, 2029	DS DP U-1402			
	8445524*PED	Sep 26, 2029				
	8609863	Jan 12, 2026	DP			
	8609863*PED	Jul 12, 2026				
	8669279	Mar 26, 2029	DP U-1402			
	8669279*PED	Sep 26, 2029				
	8791270	Jan 12, 2026	DP U-1542			
	8791270*PED	Jul 12, 2026				
	8883836	Mar 26, 2029	DP U-1402			
	8883836*PED	Sep 26, 2029				
	8895756	Jan 12, 2026	DP			
	8895756*PED	Jul 12, 2026				
	9533955	Mar 26, 2029	DP U-1949			
	9533955	Mar 26, 2029	DP U-1952			
	9533955*PED	Sep 26, 2029				
<u>BENDAMUSTINE HYDROCHLORIDE - TREANDA</u>						
N 022249	002 8436190	Oct 26, 2030	DP			
	8436190*PED	Apr 26, 2031				
	8445524	Mar 26, 2029	DS DP U-1402			
	8445524*PED	Sep 26, 2029				
	8609863	Jan 12, 2026	DP			
	8609863*PED	Jul 12, 2026				
	8669279	Mar 26, 2029	DP U-1402			
	8669279*PED	Sep 26, 2029				

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>BENDAMUSTINE HYDROCHLORIDE - TREANDA</u>						
N 022249 002	8791270	Jan 12, 2026	DP U-1542			
	8791270*PED	Jul 12, 2026				
	8883836	Mar 26, 2029	DP U-1402			
	8883836*PED	Sep 26, 2029				
	8895756	Jan 12, 2026	DP			
	8895756*PED	Jul 12, 2026				
	9533955	Mar 26, 2029	DP U-1949			
	9533955	Mar 26, 2029	DP U-1952			
	9533955*PED	Sep 26, 2029				
<u>BENDAMUSTINE HYDROCHLORIDE - TREANDA</u>						
N 022249 003	8344006	Sep 23, 2029	DP U-1402			
	8344006*PED	Mar 23, 2030				
	8445524	Mar 26, 2029	DS			
	8445524*PED	Sep 26, 2029				
	8791270	Jan 12, 2026	DP U-1542			
	8791270*PED	Jul 12, 2026				
<u>BENDAMUSTINE HYDROCHLORIDE - TREANDA</u>						
N 022249 004	8344006	Sep 23, 2029	DP U-1402			
	8344006*PED	Mar 23, 2030				
	8445524	Mar 26, 2029	DS			
	8445524*PED	Sep 26, 2029				
	8791270	Jan 12, 2026	DP U-1542			
	8791270*PED	Jul 12, 2026				
<u>BENDAMUSTINE HYDROCHLORIDE - BELRAPZO</u>						
N 205580 001	10010533	Jan 28, 2031	DP			
	11103483	Jan 28, 2031	DP U-1971			
	11103483	Jan 28, 2031	DP U-1972			
	11844783	Jan 28, 2031	U-1542			
	11844783	Jan 28, 2031	U-1971			
	11844783	Jan 28, 2031	U-1972			
	11872214	Jan 28, 2031	DP			
	8609707	Aug 11, 2031	DP U-1971			
	8609707	Aug 11, 2031	DP U-1972			
	8791270	Jan 12, 2026	DP U-1971			
	8791270	Jan 12, 2026	DP U-1972			
	9265831	Jan 28, 2031	DP			
	9572796	Jan 28, 2031	DP U-1971			
	9572796	Jan 28, 2031	DP U-1972			
	9572797	Jan 28, 2031	U-1971			
	9572797	Jan 28, 2031	U-1972			
<u>BENDAMUSTINE HYDROCHLORIDE - BENDEKA</u>						
N 208194 001	10010533	Jan 28, 2031	DP			
	10052385	Mar 15, 2033	U-1971			
	10052385	Mar 15, 2033	U-1972			
	11103483	Jan 28, 2031	DP U-1971			
	11103483	Jan 28, 2031	DP U-1972			
	11844783	Jan 28, 2031	U-1542			
	11844783	Jan 28, 2031	U-1971			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>BENDAMUSTINE HYDROCHLORIDE - BENDEKA</u>						
N 208194	001 11844783	Jan 28, 2031	U-1972			
	11872214	Jan 28, 2031	DP			
	8609707	Aug 11, 2031	DP U-1542			
	8791270	Jan 12, 2026	DP U-1790			
	9000021	Mar 15, 2033	U-1542			
	9034908	Mar 15, 2033	U-1542			
	9144568	Mar 15, 2033	U-1542			
	9265831	Jan 28, 2031	DP			
	9572796	Jan 28, 2031	DP U-1971			
	9572796	Jan 28, 2031	DP U-1972			
	9572797	Jan 28, 2031	U-1971			
	9572797	Jan 28, 2031	U-1972			
	9572887	Mar 15, 2033	U-1971			
	9572887	Mar 15, 2033	U-1972			
	9579384	Mar 15, 2033	U-1971			
	9579384	Mar 15, 2033	U-1972			
	9597397	Mar 15, 2033	U-1971			
	9597397	Mar 15, 2033	U-1972			
	9597398	Mar 15, 2033	U-1971			
	9597399	Mar 15, 2033	U-1971			
	9597399	Mar 15, 2033	U-1972			
<u>BENDAMUSTINE HYDROCHLORIDE - VIVIMUSTA</u>						
N 212209	001 11844784	Jul 29, 2042	DP			
<u>BENOXINATE HYDROCHLORIDE; FLUORESCCEIN SODIUM - FLUORESCCEIN SODIUM AND BENOXINATE HYDROCHLORIDE</u>						
N 211039	001 10293047	Nov 15, 2037	DP U-2755			
	10632197	Nov 15, 2037	DP U-2755			
	10842872	Nov 15, 2037	U-3001			
<u>BENZGALANTAMINE GLUCONATE - ZUNVEYL</u>						
N 218549	001 11795176	Jan 13, 2042	DS U-713			
	9763953	Dec 01, 2026	U-713			
<u>BENZGALANTAMINE GLUCONATE - ZUNVEYL</u>						
N 218549	002 11795176	Jan 13, 2042	DS U-713			
	9763953	Dec 01, 2026	U-713			
<u>BENZGALANTAMINE GLUCONATE - ZUNVEYL</u>						
N 218549	003 11795176	Jan 13, 2042	DS U-713			
	9763953	Dec 01, 2026	U-713			
<u>BENZOYL PEROXIDE - EPSOLAY</u>						
N 214510	001 10933046	Feb 19, 2040	DP U-3357		NP	Apr 22, 2025
	10945987	Feb 19, 2040	U-3356			
	11426378	Aug 18, 2040	U-3356			
	11541026	Feb 19, 2040	U-3356			
	11628155	Dec 27, 2040	U-3356			
	11865100	Feb 19, 2040	U-3357			
	11877997	Feb 19, 2040	U-3356			
	11986456	Feb 19, 2040	U-3947			
	12156946	Feb 03, 2028	DP			
	9687465	Nov 27, 2032	DP U-3356			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>BENZOYL PEROXIDE - EPSOLAY</u>						
N 214510	001 9868103	Aug 08, 2028	DP U-3356			
<u>BENZOYL PEROXIDE; CLINDAMYCIN PHOSPHATE - ACANYA</u>						
N 050819	001 10220049	Jun 03, 2029	DP U-916			
	10624918	Jun 03, 2029	U-916			
	8288434	Aug 05, 2029	DP U-124			
	8663699	Jun 03, 2029	U-124			
	8895070	Jun 03, 2029	U-124			
	9078870	Jun 03, 2029	DP			
<u>BENZOYL PEROXIDE; CLINDAMYCIN PHOSPHATE - ONEXTON</u>						
N 050819	002 10137142	Jun 03, 2029	DP U-916			
	10220049	Jun 03, 2029	DP U-916			
	8288434	Aug 05, 2029	DP U-1033			
	8288434	Aug 05, 2029	DP U-124			
	8288434	Aug 05, 2029	DP U-134			
	8288434	Aug 05, 2029	DP U-818			
	8288434	Aug 05, 2029	DP U-916			
	8288434	Aug 05, 2029	DP U-921			
	9504704	Jun 03, 2029	DP U-124			
	9504704	Jun 03, 2029	DP U-134			
	9504704	Jun 03, 2029	DP U-818			
	9504704	Jun 03, 2029	DP U-916			
	9561208	Jun 03, 2029	DP U-916			
<u>BENZOYL PEROXIDE; TRETINOIN - TWYNEO</u>						
N 214902	001 10420743	Jul 12, 2038	U-3194			
	10653899	Dec 30, 2030	DP U-3194			
	11071878	Dec 30, 2030	DP			
	12053546	Jun 29, 2032	DP			
	12070629	Dec 30, 2030	DP U-3987			
	12133919	May 23, 2041	DP U-3987			
	12156946	Feb 03, 2028	DP			
	8617580	Feb 03, 2028	DP			
	9868103	Aug 08, 2028	DP U-3194			
<u>BEPOTASTINE BESILATE - BEPREVE</u>						
N 022288	001 8784789	Jan 13, 2025	DP			
<u>BERDAZIMER SODIUM - ZELSUVMI</u>						
N 217424	001 10258564	Nov 22, 2034	U-3797		NCE	Jan 05, 2029
	10258564	Nov 22, 2034	U-3798			
	10258564	Nov 22, 2034	U-3799			
	10265334	Jul 03, 2032	DP			
	10322081	Jul 10, 2035	U-3793			
	10322081	Jul 10, 2035	U-3794			
	10322081	Jul 10, 2035	U-3795			
	10322081	Jul 10, 2035	U-3796			
	10376538	Aug 20, 2030	DP			
	10736839	Jul 10, 2035	U-3790			
	10736839	Jul 10, 2035	U-3791			
	10736839	Jul 10, 2035	U-3792			
	11040006	Jul 10, 2035	DP U-3789			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>BERDAZIMER SODIUM - ZELSUVMI</u>						
N 217424	001	11285098	Feb 28, 2034	DP		
		11723858	Jul 10, 2035	DP	U-3788	
		8282967	May 30, 2026	DS		
		8956658	May 30, 2026	DS		
		9289442	Jul 03, 2032	DP	U-3803	
		9526738	Sep 03, 2031	DP		
		9737561	Aug 20, 2030		U-3802	
		9855211	Feb 27, 2034	DP	U-3800	
		9855211	Feb 27, 2034	DP	U-3801	
<u>BEROTRALSTAT HYDROCHLORIDE - ORLADEYO</u>						
N 214094	001	10125102	Apr 07, 2035	DS	U-3010	NCE Dec 03, 2025
		10329260	Mar 09, 2035	DS		ODE-333 Dec 03, 2027
		10662160	Nov 01, 2039	DS	U-3010	
		10689346	Mar 09, 2035		U-3010	
		11117867	Nov 01, 2039	DP	U-3010	
		11230530	Mar 09, 2035		U-3300	
		11618733	Nov 01, 2039		U-3300	
		11708333	Mar 09, 2035		U-3300	
		12116346	Mar 09, 2035	DS		
<u>BEROTRALSTAT HYDROCHLORIDE - ORLADEYO</u>						
N 214094	002	10125102	Apr 07, 2035	DS	U-3010	NCE Dec 03, 2025
		10329260	Mar 09, 2035	DS		ODE-333 Dec 03, 2027
		10662160	Nov 01, 2039	DS	U-3010	
		10689346	Mar 09, 2035		U-3010	
		11117867	Nov 01, 2039	DP	U-3010	
		11230530	Mar 09, 2035		U-3300	
		11618733	Nov 01, 2039		U-3300	
		11708333	Mar 09, 2035		U-3300	
		12116346	Mar 09, 2035	DS		
<u>BESIFLOXACIN HYDROCHLORIDE - BESIVANCE</u>						
N 022308	001	8415342	Nov 07, 2030		U-80	
		8481526	Jan 09, 2031	DS		
		8604020	Mar 12, 2030	DP		
		8937062	Nov 13, 2029		U-80	
<u>BETAMETHASONE DIPROPIONATE - SERNIVO</u>						
N 208079	001	10179137	Aug 31, 2030	DP	U-1858	
		9364485	Aug 31, 2030	DP	U-1858	
		9433630	Aug 31, 2030	DP	U-1858	
		9439911	Aug 31, 2030	DP	U-1858	
		9655907	Aug 31, 2030	DP	U-1858	
		9775851	Aug 31, 2030	DP	U-1858	
		9877974	Aug 31, 2030	DP	U-1858	
<u>BETAMETHASONE DIPROPIONATE; CALCIPOTRIENE - ENSTILAR</u>						
N 207589	001	10130640	Jun 10, 2031	DP		
		10130640*PED	Dec 10, 2031			
		10617698	Jun 10, 2031	DP		
		10660908	Jun 10, 2031	DP	U-2627	
		10682364	Jun 10, 2031	DP		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>BETAMETHASONE DIPROPIONATE; CALCIPOTRIENE - ENSTILAR</u>						
N 207589	001	10688108	Jun 10, 2031	U-2627		
		10716799	Jun 10, 2031	DP		
		9119781	Jun 10, 2031	DP U-1761		
		9119781	Jun 10, 2031	DP U-2627		
		9119781*PED	Dec 10, 2031			
		9566286	Jun 10, 2031	DP		
<u>BETAMETHASONE DIPROPIONATE; CALCIPOTRIENE - WYNZORA</u>						
N 213422	001	10265265	Sep 27, 2027	DP		
		11696919	Mar 18, 2039	DP		
<u>BETRIXABAN - BEVYXXA</u>						
N 208383	001	7598276	Nov 08, 2026	DS		
		8404724	Mar 29, 2031	DP U-2034		
		8557852	Sep 08, 2028	U-1167		
		8557852	Sep 08, 2028	U-2030		
		8987463	Dec 28, 2030	DP		
		9555023	Nov 07, 2026	U-1502		
<u>BETRIXABAN - BEVYXXA</u>						
N 208383	002	7598276	Nov 08, 2026	DS		
		8404724	Mar 29, 2031	DP U-2034		
		8557852	Sep 08, 2028	U-1167		
		8557852	Sep 08, 2028	U-2030		
		8987463	Dec 28, 2030	DP		
		9555023	Nov 07, 2026	U-1502		
<u>BEXAGLIFLOZIN - BRENZAVVY</u>						
N 214373	001	10533032	Jul 03, 2031	U-2214	NCE	Jan 20, 2028
		10981942	Jun 13, 2031	DS DP		
		7838499	Jan 30, 2029	DS DP U-2214		
		8106021	Aug 22, 2028	U-2214		
		8802637	Aug 22, 2028	DS DP U-2214		
		8987323	May 14, 2032	DS DP		
<u>BICTEGRAVIR SODIUM; EMTRICITABINE; TENOFOVIR ALAFENAMIDE FUMARATE - BIKTARVY</u>						
N 210251	001	10385067	Jun 19, 2035	U-257	I-942	Feb 23, 2027
		10548846	Nov 08, 2036	DP	M-305	Apr 24, 2027
		11744802	Nov 08, 2036	DP	ODE-256	Jun 18, 2026
		7390791	Apr 17, 2025	DS DP	ODE-468	Feb 23, 2031
		7390791*PED	Oct 17, 2025			
		8754065	Aug 15, 2032	DS DP U-257		
		8754065*PED	Feb 15, 2033			
		9216996	Dec 19, 2033	DS DP		
		9296769	Aug 15, 2032	DS DP U-257		
		9296769*PED	Feb 15, 2033			
		9708342	Jun 19, 2035	DS DP		
		9732092	Dec 19, 2033	DS DP		
<u>BICTEGRAVIR SODIUM; EMTRICITABINE; TENOFOVIR ALAFENAMIDE FUMARATE - BIKTARVY</u>						
N 210251	002	10385067	Jun 19, 2035	U-257	I-942	Feb 23, 2027
		7390791	Apr 17, 2025	DS DP	ODE-378	Oct 07, 2028
		7390791*PED	Oct 17, 2025		ODE-468	Feb 23, 2031

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>BICTEGRAVIR SODIUM; EMTRICITABINE; TENOFOVIR ALAFENAMIDE FUMARATE - BIKTARVY</u>						
N 210251	002 8754065	Aug 15, 2032	DS DP U-257			
	8754065*PED	Feb 15, 2033				
	9216996	Dec 19, 2033	DS DP			
	9296769	Aug 15, 2032	DS DP U-257			
	9296769*PED	Feb 15, 2033				
	9708342	Jun 19, 2035	DS DP			
	9732092	Dec 19, 2033	DS DP			
<u>BIMATOPROST - LUMIGAN</u>						
N 022184	001 7851504	Jun 13, 2027	DS DP			
	8278353	Mar 16, 2025	DP			
	8299118	Mar 16, 2025		U-1295		
	8309605	Mar 16, 2025		U-1293		
	8309605	Mar 16, 2025		U-1294		
	8338479	Mar 16, 2025	DP U-1295			
	8524777	Mar 16, 2025		U-1235		
	8586630	Mar 16, 2025		U-1458		
	8772338	Mar 16, 2025	DP U-1528			
	8933120	Mar 16, 2025	DP			
	8933127	Mar 16, 2025	DP			
	9155716	Mar 16, 2025	DP U-1528			
	9241918	Mar 16, 2025	DP U-1814			
<u>BIMATOPROST - DURYSTA</u>						
N 211911	001 10441543	Dec 19, 2026	DP			
	7799336	Apr 24, 2029	DP			
	8206737	Apr 07, 2027		U-2759		
	8629185	Jul 15, 2031	DS DP			
	8673341	Feb 19, 2025		U-2759		
	9149428	Dec 19, 2026	DP			
	9492316	Oct 31, 2034	DP			
	9980974	Oct 31, 2034		U-2759		
<u>BINIMETINIB - MEKTOVI</u>						
N 210498	001 10005761	Aug 27, 2030		U-2331	I-928	Oct 11, 2026
	10005761	Aug 27, 2030		U-3737	ODE-194	Jun 27, 2025
	7777050	Mar 13, 2025	DS DP			
	9314464	Jul 04, 2031		U-2332		
	9314464	Jul 04, 2031		U-3737		
	9562016	Oct 18, 2033	DS DP			
	9598376	Oct 18, 2033		U-2330		
	9850229	Aug 27, 2030		U-2333		
	9980944	Oct 18, 2033		U-2334		
<u>BIRCH TRITERPENES - FILSUVUZ</u>						
N 215064	001 11083733	Jan 04, 2039	DP		NCE	Dec 18, 2028
	11266660	Jan 04, 2039		U-3811	ODE-460	Dec 18, 2030
	8828444	Jun 21, 2025	DP			
	9352041	Nov 24, 2030		U-3811		
	9827214	Nov 24, 2030		U-3811		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>BIVALIRUDIN - ANGIOMAX</u>						
N 020873	001 7582727	Jul 27, 2028	DP			
	7598343	Jul 27, 2028	DP			
<u>BIVALIRUDIN - ANGIOMAX RTU</u>						
N 211215	001 11903993	May 20, 2039	U-3817			
	11918622	May 20, 2039	U-3817			
	11992514	May 20, 2039	DP			
<u>BOCEPREVIR - VICTRELIS</u>						
N 202258	001 7772178	Nov 11, 2027	DP U-1128			
	8119602	Mar 17, 2027	U-1233			
<u>BORTEZOMIB - BORTEZOMIB</u>						
N 205004	001 8962572	Nov 03, 2032	DP			
<u>BORTEZOMIB - BORTEZOMIB</u>						
N 215331	001 11679119	Sep 23, 2042	U-3632			
	11679119	Sep 23, 2042	U-3633			
	11752164	Sep 23, 2042	U-3632			
	11752164	Sep 23, 2042	U-3633			
	12005069	Sep 23, 2042	DP			
<u>BORTEZOMIB - BORTEZOMIB</u>						
N 215331	002 11679119	Sep 23, 2042	U-3632			
	11679119	Sep 23, 2042	U-3633			
	11752164	Sep 23, 2042	U-3632			
	11752164	Sep 23, 2042	U-3633			
	12005069	Sep 23, 2042	DP			
<u>BOSENTAN - TRACLEER</u>						
N 209279	001 7959945	Dec 28, 2027	DP			
	8309126	May 15, 2026	DP			
<u>BOSUTINIB MONOHYDRATE - BOSULIF</u>						
N 203341	001 11103497	Feb 28, 2034	U-3216		I-923	Sep 26, 2026
	11103497	Feb 28, 2034	U-3217		ODE-163	Dec 19, 2024
	11103497*PED	Aug 28, 2034			ODE-444	Sep 26, 2030
	7417148	Dec 11, 2025	U-1283		PED	Jun 19, 2025
	7417148	Dec 11, 2025	U-3707		PED	Mar 26, 2027
	7417148	Dec 11, 2025	U-3708		PED	Mar 26, 2031
	7417148*PED	Jun 11, 2026				
	7767678	Nov 23, 2026	DS DP			
	7767678*PED	May 23, 2027				
	7919625	Dec 11, 2025	DP			
	7919625*PED	Jun 11, 2026				
<u>BOSUTINIB MONOHYDRATE - BOSULIF</u>						
N 203341	002 11103497	Feb 28, 2034	U-3216		I-923	Sep 26, 2026
	11103497	Feb 28, 2034	U-3217		ODE-163	Dec 19, 2024
	11103497*PED	Aug 28, 2034			ODE-444	Sep 26, 2030
	7417148	Dec 11, 2025	U-1283		PED	Jun 19, 2025
	7417148	Dec 11, 2025	U-3707		PED	Mar 26, 2027
	7417148	Dec 11, 2025	U-3708		PED	Mar 26, 2031
	7417148*PED	Jun 11, 2026				

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>BOSUTINIB MONOHYDRATE - BOSULIF</u>						
N 203341 002	7767678	Nov 23, 2026	DS DP			
	7767678*PED	May 23, 2027				
	7919625	Dec 11, 2025	DP			
	7919625*PED	Jun 11, 2026				
<u>BOSUTINIB MONOHYDRATE - BOSULIF</u>						
N 203341 003	11103497	Feb 28, 2034	U-3216		I-923	Sep 26, 2026
	11103497	Feb 28, 2034	U-3217		ODE-163	Dec 19, 2024
	11103497*PED	Aug 28, 2034			ODE-444	Sep 26, 2030
	7417148	Dec 11, 2025	U-1283		PED	Jun 19, 2025
	7417148	Dec 11, 2025	U-3707		PED	Mar 26, 2027
	7417148	Dec 11, 2025	U-3708		PED	Mar 26, 2031
	7417148*PED	Jun 11, 2026				
	7767678	Nov 23, 2026	DS DP			
	7767678*PED	May 23, 2027				
	7919625	Dec 11, 2025	DP			
	7919625*PED	Jun 11, 2026				
<u>BOSUTINIB MONOHYDRATE - BOSULIF</u>						
N 217729 001	11103497	Feb 28, 2034	U-3216		NP	Sep 26, 2026
	11103497	Feb 28, 2034	U-3217		ODE-444	Sep 26, 2030
	11103497*PED	Aug 28, 2034			PED	Mar 26, 2027
	7417148	Dec 11, 2025	U-3707		PED	Mar 26, 2031
	7417148	Dec 11, 2025	U-3708			
	7417148*PED	Jun 11, 2026				
	7767678	Nov 23, 2026	DS DP			
	7767678*PED	May 23, 2027				
	7919625	Dec 11, 2025	DP			
	7919625*PED	Jun 11, 2026				
<u>BOSUTINIB MONOHYDRATE - BOSULIF</u>						
N 217729 002	11103497	Feb 28, 2034	U-3216		NP	Sep 26, 2026
	11103497	Feb 28, 2034	U-3217		ODE-444	Sep 26, 2030
	11103497*PED	Aug 28, 2034			PED	Mar 26, 2027
	7417148	Dec 11, 2025	U-3707		PED	Mar 26, 2031
	7417148	Dec 11, 2025	U-3708			
	7417148*PED	Jun 11, 2026				
	7767678	Nov 23, 2026	DS DP			
	7767678*PED	May 23, 2027				
	7919625	Dec 11, 2025	DP			
	7919625*PED	Jun 11, 2026				
<u>BREMELANOTIDE ACETATE - VYLEESI (AUTOINJECTOR)</u>						
N 210557 001	10286034	Nov 05, 2033	U-2568			
	11590209	Apr 29, 2041	U-3539			
	6794489	Jun 28, 2025	DS DP			
	9352013	Nov 05, 2033	U-2568			
	9700592	Nov 05, 2033	U-2568			
<u>BREXANOLONE - ZULRESSO</u>						
N 211371 001	10117951	Mar 13, 2029	DP		NPP	Jun 16, 2025
	10251894	Nov 27, 2033	U-2552			
	10322139	Jan 23, 2033	DP			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>BREXANOLONE - ZULRESSO</u>						
N 211371 001	10940156	Mar 08, 2037	U-2552			
	7635773	Mar 13, 2029	DP			
	8410077	Mar 13, 2029	DP			
	9200088	Mar 13, 2029	DP			
	9750822	Mar 13, 2029	DP			
<u>BREXPIPIRAZOLE - REXULTI</u>						
N 205422 001	10307419	Oct 12, 2032	DP		I-913	May 10, 2026
	10307419*PED	Apr 12, 2033			M-14	May 08, 2027
	7888362	Apr 12, 2026	DS	Y	PED	Nov 10, 2026
	7888362*PED	Oct 12, 2026			PED	Nov 08, 2027
	8349840	Apr 12, 2026	DP U-1529			
	8349840*PED	Oct 12, 2026				
	8618109	Apr 12, 2026	U-3281			
	8618109	Apr 12, 2026	U-543			
	8618109*PED	Oct 12, 2026				
	9839637	Apr 12, 2026	DP U-1529			
	9839637	Apr 12, 2026	DP U-3281			
	9839637	Apr 12, 2026	DP U-543			
	9839637*PED	Oct 12, 2026				
	RE48059	Dec 23, 2028	DS			
	RE48059*PED	Jun 23, 2029				
<u>BREXPIPIRAZOLE - REXULTI</u>						
N 205422 002	10307419	Oct 12, 2032	DP		I-913	May 10, 2026
	10307419*PED	Apr 12, 2033			M-14	May 08, 2027
	7888362	Apr 12, 2026	DS	Y	PED	Nov 10, 2026
	7888362*PED	Oct 12, 2026			PED	Nov 08, 2027
	8349840	Apr 12, 2026	DP U-1529			
	8349840*PED	Oct 12, 2026				
	8618109	Apr 12, 2026	U-3281			
	8618109	Apr 12, 2026	U-543			
	8618109*PED	Oct 12, 2026				
	9839637	Apr 12, 2026	DP U-1529			
	9839637	Apr 12, 2026	DP U-3281			
	9839637	Apr 12, 2026	DP U-543			
	9839637*PED	Oct 12, 2026				
	RE48059	Dec 23, 2028	DS			
	RE48059*PED	Jun 23, 2029				
<u>BREXPIPIRAZOLE - REXULTI</u>						
N 205422 003	10307419	Oct 12, 2032	DP		I-913	May 10, 2026
	10307419*PED	Apr 12, 2033			M-14	May 08, 2027
	7888362	Apr 12, 2026	DS	Y	PED	Nov 10, 2026
	7888362*PED	Oct 12, 2026			PED	Nov 08, 2027
	8349840	Apr 12, 2026	DP U-1529			
	8349840*PED	Oct 12, 2026				
	8618109	Apr 12, 2026	U-3281			
	8618109	Apr 12, 2026	U-543			
	8618109*PED	Oct 12, 2026				
	9839637	Apr 12, 2026	DP U-1529			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>BREXPIRAZOLE - REXULTI</u>						
N 205422 003	9839637	Apr 12, 2026	DP U-3281			
	9839637	Apr 12, 2026	DP U-543			
	9839637*PED	Oct 12, 2026				
	RE48059	Dec 23, 2028	DS			
	RE48059*PED	Jun 23, 2029				
<u>BREXPIRAZOLE - REXULTI</u>						
N 205422 004	10307419	Oct 12, 2032	DP		I-913	May 10, 2026
	10307419*PED	Apr 12, 2033			M-14	May 08, 2027
	7888362	Apr 12, 2026	DS	Y	PED	Nov 10, 2026
	7888362*PED	Oct 12, 2026			PED	Nov 08, 2027
	8349840	Apr 12, 2026	DP U-1529			
	8349840*PED	Oct 12, 2026				
	8618109	Apr 12, 2026	U-3281			
	8618109	Apr 12, 2026	U-543			
	8618109*PED	Oct 12, 2026				
	9839637	Apr 12, 2026	DP U-1529			
	9839637	Apr 12, 2026	DP U-3281			
	9839637	Apr 12, 2026	DP U-543			
	9839637*PED	Oct 12, 2026				
	RE48059	Dec 23, 2028	DS			
	RE48059*PED	Jun 23, 2029				
<u>BREXPIRAZOLE - REXULTI</u>						
N 205422 005	10307419	Oct 12, 2032	DP		I-913	May 10, 2026
	10307419*PED	Apr 12, 2033			M-14	May 08, 2027
	7888362	Apr 12, 2026	DS	Y	PED	Nov 10, 2026
	7888362*PED	Oct 12, 2026			PED	Nov 08, 2027
	8349840	Apr 12, 2026	DP U-1529			
	8349840*PED	Oct 12, 2026				
	8618109	Apr 12, 2026	U-3281			
	8618109	Apr 12, 2026	U-543			
	8618109*PED	Oct 12, 2026				
	9839637	Apr 12, 2026	DP U-1529			
	9839637	Apr 12, 2026	DP U-3281			
	9839637	Apr 12, 2026	DP U-543			
	9839637*PED	Oct 12, 2026				
	RE48059	Dec 23, 2028	DS			
	RE48059*PED	Jun 23, 2029				
<u>BREXPIRAZOLE - REXULTI</u>						
N 205422 006	7888362	Apr 12, 2026	DS	Y	I-913	May 10, 2026
	7888362*PED	Oct 12, 2026			M-14	May 08, 2027
	8349840	Apr 12, 2026	DP U-1529		PED	Nov 10, 2026
	8349840*PED	Oct 12, 2026			PED	Nov 08, 2027
	8618109	Apr 12, 2026	U-3281			
	8618109	Apr 12, 2026	U-543			
	8618109*PED	Oct 12, 2026				
	9839637	Apr 12, 2026	DP U-1529			
	9839637	Apr 12, 2026	DP U-3281			
	9839637	Apr 12, 2026	DP U-543			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>BREXPIRAZOLE - REXULTI</u>						
N 205422	006 9839637*PED	Oct 12, 2026				
	RE48059	Dec 23, 2028	DS			
	RE48059*PED	Jun 23, 2029				
<u>BRIGATINIB - ALUNBRIG</u>						
N 208772	001 10385078	Nov 10, 2035	DS DP U-2837		ODE-300	May 22, 2027
	9012462	Apr 28, 2031	DS			
	9273077	May 21, 2029		U-2837		
	9611283	Apr 10, 2034		U-2837		
<u>BRIGATINIB - ALUNBRIG</u>						
N 208772	002 10385078	Nov 10, 2035	DS DP U-2837		ODE-300	May 22, 2027
	9012462	Apr 28, 2031	DS			
	9273077	May 21, 2029		U-2837		
	9611283	Apr 10, 2034		U-2837		
<u>BRIGATINIB - ALUNBRIG</u>						
N 208772	003 10385078	Nov 10, 2035	DS DP U-2837		ODE-300	May 22, 2027
	9012462	Apr 28, 2031	DS			
	9273077	May 21, 2029		U-2837		
	9611283	Apr 10, 2034		U-2837		
<u>BRILLIANT BLUE G - TISSUEBLUE</u>						
N 209569	001				ODE-282	Dec 20, 2026
<u>BRIMONIDINE TARTRATE - QOLIANA</u>						
N 021764	001 7265117	Aug 19, 2025	DP			
<u>BRIMONIDINE TARTRATE - MIRVASO</u>						
N 204708	001 10201517	Jun 13, 2031	DP			
	7439241	Aug 25, 2025		U-1428		
	8053427	Jun 13, 2031	DP U-1428			
	8163725	Jun 13, 2031	DP			
	8231885	May 24, 2025	DP			
	8410102	May 24, 2025		U-1428		
	8426410	May 24, 2025		U-1428		
	8513247	Mar 25, 2031	DP U-1428			
	8513249	Mar 25, 2031	DP U-1428			
	9861631	Mar 25, 2031		U-1428		
	9861632	Mar 25, 2031		U-1428		
<u>BRIMONIDINE TARTRATE - LUMIFY</u>						
N 208144	001 11596600	Jul 27, 2029		U-2222		
	11833245	Jul 27, 2029		U-2222		
	8293742	Jul 14, 2030		U-2222		
	9259425	Jul 14, 2030		U-2222		
<u>BRIMONIDINE TARTRATE - LUMIFY PRESERVATIVE FREE</u>						
N 218424	001 11596600	Jul 27, 2029		U-2222		
	8293742	Jul 14, 2030		U-2222		
	9259425	Jul 14, 2030		U-2222		
<u>BRIMONIDINE TARTRATE; BRINZOLAMIDE - SIMBRINZA</u>						
N 204251	001 9044484	Oct 30, 2030	DP			
	9421265	Jun 17, 2030	DP			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>BRINCIDOFOVIR - TEMBEXA</u>						
N 214460	001 9303051	Aug 31, 2031	DS DP U-3165		ODE-354	Jun 04, 2028
<u>BRINCIDOFOVIR - TEMBEXA</u>						
N 214461	001 10112909	Oct 10, 2034	U-3165		ODE-354	Jun 04, 2028
	10487061	Oct 10, 2034	DP U-3165			
	8962829	Oct 10, 2034	DS DP			
	9303051	Aug 31, 2031	DS DP U-3165			
	9371344	Oct 10, 2034	DP			
<u>BRIVARACETAM - BRIVIACT</u>						
N 205836	001 10729653	Apr 09, 2030	DP			
	6911461	Feb 21, 2026	DS DP U-2295			
<u>BRIVARACETAM - BRIVIACT</u>						
N 205836	002 10729653	Apr 09, 2030	DP			
	6911461	Feb 21, 2026	DS DP U-2295			
<u>BRIVARACETAM - BRIVIACT</u>						
N 205836	003 10729653	Apr 09, 2030	DP			
	6911461	Feb 21, 2026	DS DP U-2295			
<u>BRIVARACETAM - BRIVIACT</u>						
N 205836	004 10729653	Apr 09, 2030	DP			
	6911461	Feb 21, 2026	DS DP U-2295			
<u>BRIVARACETAM - BRIVIACT</u>						
N 205836	005 10729653	Apr 09, 2030	DP			
	6911461	Feb 21, 2026	DS DP U-2295			
<u>BRIVARACETAM - BRIVIACT</u>						
N 205837	001 6911461	Feb 21, 2026	DS DP U-1815			
	6911461	Feb 21, 2026	DS DP U-2130			
<u>BRIVARACETAM - BRIVIACT</u>						
N 205838	001 6911461	Feb 21, 2026	DS DP U-2295			
<u>BROMFENAC SODIUM - PROLENSA</u>						
N 203168	001 10085958	Nov 19, 2032	DP			
	8129431	Sep 11, 2025	DS DP			
	9517220	Nov 11, 2033	U-1933			
<u>BROMFENAC SODIUM - BROMSITE</u>						
N 206911	001 RE50218	Mar 05, 2029	DP U-1834			
<u>BROMOCRIPTINE MESYLATE - CYCLOSET</u>						
N 020866	001 10688094	Apr 30, 2032	U-2870			
	10688094	Apr 30, 2032	U-2871			
	10688094	Apr 30, 2032	U-2872			
	10688094	Apr 30, 2032	U-2873			
	10688094	Apr 30, 2032	U-2874			
	10688094	Apr 30, 2032	U-2875			
	10688094	Apr 30, 2032	U-2876			
	10688094	Apr 30, 2032	U-2877			
	10688094	Apr 30, 2032	U-2878			
	10688094	Apr 30, 2032	U-2879			
	10688094	Apr 30, 2032	U-2880			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>BROMOCRIPTINE MESYLATE - CYCLOSET</u>						
N 020866 001	10688094	Apr 30, 2032	U-2881			
	10688094	Apr 30, 2032	U-2882			
	10688094	Apr 30, 2032	U-2883			
	10688094	Apr 30, 2032	U-2884			
	10688094	Apr 30, 2032	U-2885			
	10688094	Apr 30, 2032	U-2886			
	10688094	Apr 30, 2032	U-2887			
	10688094	Apr 30, 2032	U-2888			
	10688155	Jun 07, 2030	U-2281			
	10688155	Jun 07, 2030	U-2890			
	10688155	Jun 07, 2030	U-2891			
	10688155	Jun 07, 2030	U-2892			
	10688155	Jun 07, 2030	U-2893			
	10688155	Jun 07, 2030	U-2894			
	10688155	Jun 07, 2030	U-2895			
	10688155	Jun 07, 2030	U-2896			
	10688155	Jun 07, 2030	U-2897			
	10688155	Jun 07, 2030	U-2898			
	10688155	Jun 07, 2030	U-2899			
	10688155	Jun 07, 2030	U-2900			
	10688155	Jun 07, 2030	U-2901			
	10688155	Jun 07, 2030	U-2902			
	10688155	Jun 07, 2030	U-2903			
	10688155	Jun 07, 2030	U-2904			
	10688155	Jun 07, 2030	U-2905			
	10688155	Jun 07, 2030	U-2906			
	10688155	Jun 07, 2030	U-2907			
	10688155	Jun 07, 2030	U-2908			
	10688155	Jun 07, 2030	U-2909			
	10688155	Jun 07, 2030	U-2910			
	10688155	Jun 07, 2030	U-2911			
	10688155	Jun 07, 2030	U-2912			
	10688155	Jun 07, 2030	U-2913			
	10688155	Jun 07, 2030	U-2914			
	10688155	Jun 07, 2030	U-2915			
	10688155	Jun 07, 2030	U-2916			
	10688155	Jun 07, 2030	U-2917			
	10688155	Jun 07, 2030	U-2918			
	10688155	Jun 07, 2030	U-2919			
	10688155	Jun 07, 2030	U-2920			
	10688155	Jun 07, 2030	U-2921			
	10688155	Jun 07, 2030	U-2922			
	10688155	Jun 07, 2030	U-2923			
	10688155	Jun 07, 2030	U-2924			
	10688155	Jun 07, 2030	U-2925			
	10688155	Jun 07, 2030	U-2926			
	10688155	Jun 07, 2030	U-2927			
	10688155	Jun 07, 2030	U-2928			
	10688155	Jun 07, 2030	U-2929			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>BROMOCRIPTINE MESYLATE - CYCLOSET</u>						
N 020866 001	10688155	Jun 07, 2030	U-2930			
	10688155	Jun 07, 2030	U-2931			
	10688155	Jun 07, 2030	U-2932			
	10688155	Jun 07, 2030	U-2933			
	10688155	Jun 07, 2030	U-2934			
	10688155	Jun 07, 2030	U-2935			
	10688155	Jun 07, 2030	U-2936			
	10688155	Jun 07, 2030	U-2937			
	11000522	Apr 30, 2032	U-3119			
	11000522	Apr 30, 2032	U-3120			
	11000522	Apr 30, 2032	U-3121			
	11000522	Apr 30, 2032	U-3122			
	11666567	Apr 30, 2032	U-4015			
	11666567	Apr 30, 2032	U-4016			
	11666567	Apr 30, 2032	U-4017			
	11666567	Apr 30, 2032	U-4018			
	8431155	Apr 30, 2032	DP U-976			
	8613947	Apr 30, 2032	DP U-976			
	8877708	Jun 07, 2030	DP U-1706			
	9192576	Apr 30, 2032	DP U-976			
	9352025	Jun 07, 2030	U-2111			
	9352025	Jun 07, 2030	U-2112			
	9352025	Jun 07, 2030	U-2113			
	9352025	Jun 07, 2030	U-2114			
	9352025	Jun 07, 2030	U-2115			
	9352025	Jun 07, 2030	U-2116			
	9352025	Jun 07, 2030	U-2117			
	9352025	Jun 07, 2030	U-2118			
	9352025	Jun 07, 2030	U-2119			
	9522117	Apr 30, 2032	DP U-1939			
	9522117	Apr 30, 2032	DP U-976			
	9700555	Apr 30, 2032	DP U-2183			
	9700555	Apr 30, 2032	DP U-2184			
	9700555	Apr 30, 2032	DP U-2185			
	9700555	Apr 30, 2032	DP U-2186			
	9700555	Apr 30, 2032	DP U-2187			
	9700555	Apr 30, 2032	DP U-2188			
	9700555	Apr 30, 2032	DP U-2189			
	9700555	Apr 30, 2032	DP U-2190			
	9700555	Apr 30, 2032	DP U-2191			
	9700555	Apr 30, 2032	DP U-2192			
	9700555	Apr 30, 2032	DP U-2193			
	9700555	Apr 30, 2032	DP U-2194			
	9700555	Apr 30, 2032	DP U-2195			
	9700555	Apr 30, 2032	DP U-2196			
	9700555	Apr 30, 2032	DP U-2197			
	9700555	Apr 30, 2032	DP U-2198			
	9895422	Jun 07, 2030	U-2114			
	9895422	Jun 07, 2030	U-2116			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>BROMOCRIPTINE MESYLATE - CYCLOSET</u>						
N 020866	001	9895422	Jun 07, 2030	U-2281		
		9895422	Jun 07, 2030	U-2282		
		9895422	Jun 07, 2030	U-2283		
		9895422	Jun 07, 2030	U-2284		
		9895422	Jun 07, 2030	U-2285		
		9895422	Jun 07, 2030	U-2286		
		9895422	Jun 07, 2030	U-2287		
		9993474	Apr 30, 2032	U-2384		
		9993474	Apr 30, 2032	U-2385		
		9993474	Apr 30, 2032	U-2386		
		9993474	Apr 30, 2032	U-2387		
		9993474	Apr 30, 2032	U-2388		
		9993474	Apr 30, 2032	U-2389		
		9993474	Apr 30, 2032	U-2390		
		9993474	Apr 30, 2032	U-2391		
		9993474	Apr 30, 2032	U-2392		
		9993474	Apr 30, 2032	U-2393		
<u>BUDESONIDE - UCERIS</u>						
N 203634	001	10307375	Sep 07, 2031	DP		
		10660858	Sep 07, 2031	DP		
		8895064	Sep 07, 2031	DP		
		9132093	Sep 07, 2031	DP		
		9192581	Sep 07, 2031	DP	U-1325	
<u>BUDESONIDE - ORTIKOS</u>						
N 211929	001	10172802	Sep 09, 2036	U-2554		
		9707182	Sep 09, 2036	DP	U-2554	
<u>BUDESONIDE - ORTIKOS</u>						
N 211929	002	10172802	Sep 09, 2036	U-2554		
		9707182	Sep 09, 2036	DP	U-2554	
<u>BUDESONIDE - EOHILIA</u>						
N 213976	001	10293052	Nov 22, 2028	DP	NP	Feb 09, 2027
		11197822	Nov 09, 2026	DP	ODE-466	Feb 09, 2031
		11260064	Jan 10, 2039	DP		
		11357859	Nov 12, 2028	U-3820		
		11413296	Nov 09, 2026	DP		
		11564934	Jan 10, 2039	U-3820		
		8324192	Aug 03, 2029	U-3820		
		8497258	Nov 09, 2026	U-3820		
		8679545	Nov 09, 2026	DP		
		8975243	Nov 09, 2026	U-3820		
		9050368	Aug 01, 2029	DP		
		9119863	Nov 09, 2026	U-3820		
<u>BUDESONIDE - TARPEYO</u>						
N 215935	001	11896719	Jan 23, 2043	U-3810	ODE-389	Dec 15, 2028
		8491932	May 07, 2029	DP	ODE-464	Dec 20, 2030
		8491932	May 07, 2029	DP	U-3781	

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>BUDESONIDE; FORMOTEROL FUMARATE - SYMBICORT AEROSPHERE</u>						
N 216579	001 10716753	May 28, 2030	DP U-3203			
	9415009	May 28, 2030	U-3203			
<u>BUDESONIDE; FORMOTEROL FUMARATE DIHYDRATE - SYMBICORT</u>						
N 021929	001 7587988	Apr 10, 2026	DP			
	7587988*PED	Oct 10, 2026				
	8387615	Mar 26, 2027	DP			
	8387615*PED	Sep 26, 2027				
	8528545	Oct 16, 2028	DP			
	8528545*PED	Apr 16, 2029				
	8616196	Apr 07, 2029	DP			
	8616196*PED	Oct 07, 2029				
	8875699*PED	May 10, 2025				
<u>BUDESONIDE; FORMOTEROL FUMARATE DIHYDRATE - SYMBICORT</u>						
N 021929	002 7587988	Apr 10, 2026	DP			
	7587988*PED	Oct 10, 2026				
	8387615	Mar 26, 2027	DP			
	8387615*PED	Sep 26, 2027				
	8528545	Oct 16, 2028	DP			
	8528545*PED	Apr 16, 2029				
	8616196	Apr 07, 2029	DP			
	8616196*PED	Oct 07, 2029				
	8875699*PED	May 10, 2025				
<u>BUDESONIDE; FORMOTEROL FUMARATE; GLYCOPYRROLATE - BREZTRI AEROSPHERE</u>						
N 212122	001 10716753	May 28, 2030	DP U-2889			
	11331442	Oct 05, 2038	DP			
	11833292	Oct 05, 2038	DP			
	8324266	May 28, 2030	U-2889			
	8703806	May 28, 2030	U-2889			
	8808713	May 28, 2030	DP U-2889			
	8815258	Mar 17, 2031	U-2889			
	9415009	May 28, 2030	U-2889			
	9463161	May 28, 2030	DP U-2889			
<u>BUPIVACAINE - EXPAREL</u>						
N 022496	001 11033495	Jan 22, 2041	DP U-3182		I-929	Nov 09, 2026
	11179336	Jan 22, 2041	DP U-3250			
	11278494	Jan 22, 2041	DP U-3250			
	11304904	Jan 22, 2041	DP U-3346			
	11311486	Jan 22, 2041	DP U-3250			
	11357727	Jan 22, 2041	DP U-3380			
	11426348	Jan 22, 2041	DP U-3380			
	11452691	Jan 22, 2041	DP U-3439			
	11819574	Jan 22, 2041	DP U-3250			
	11819575	Jan 22, 2041	DP U-3250			
	11918565	Feb 02, 2043	U-3841			
	11925706	Jan 22, 2041	DP U-3380			
	11931459	Mar 17, 2042	U-3839			
	11931459	Mar 17, 2042	U-3840			
	12144890	Jan 22, 2041	DP U-4033			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>BUPIVACAINE - EXPAREL</u>						
N 022496	001 12151024	Jan 22, 2041	DP U-4037			
	12156940	Jul 02, 2044	DP			
<u>BUPIVACAINE - EXPAREL</u>						
N 022496	002 11033495	Jan 22, 2041	DP U-3182		I-929	Nov 09, 2026
	11179336	Jan 22, 2041	DP U-3250			
	11278494	Jan 22, 2041	DP U-3250			
	11304904	Jan 22, 2041	DP U-3346			
	11311486	Jan 22, 2041	DP U-3250			
	11357727	Jan 22, 2041	DP U-3380			
	11426348	Jan 22, 2041	DP U-3380			
	11452691	Jan 22, 2041	DP U-3439			
	11819574	Jan 22, 2041	DP U-3250			
	11819575	Jan 22, 2041	DP U-3250			
	11918565	Feb 02, 2043	U-3841			
	11925706	Jan 22, 2041	DP U-3380			
	11931459	Mar 17, 2042	U-3839			
	11931459	Mar 17, 2042	U-3840			
	12144890	Jan 22, 2041	DP U-4033			
	12151024	Jan 22, 2041	DP U-4037			
	12156940	Jul 02, 2044	DP			
<u>BUPIVACAINE - POSIMIR</u>						
N 204803	001 11400019	Jan 12, 2041	DP			
	11771624	Jan 12, 2041	U-3724			
	8153149	Sep 15, 2025	DP			
	8153661	Sep 15, 2025	U-3074			
	8753665	Sep 15, 2025	DP U-3074			
	8846072	Sep 15, 2025	DP U-3074			
<u>BUPIVACAINE HYDROCHLORIDE - XARACOLL</u>						
N 209511	001 11746141	Jan 09, 2033	DP			
	RE47826	May 20, 2029	U-2949			
<u>BUPIVACAINE; MELOXICAM - ZYNRELEF KIT</u>						
N 211988	001 10098957	Apr 20, 2035	U-3118		I-933	Jan 23, 2027
	10213510	Apr 20, 2035	DP U-3118			
	10398686	Mar 13, 2034	DP			
	10632199	Apr 20, 2035	DP U-3118			
	10898575	Apr 20, 2035	DP U-3118			
	10980886	Apr 20, 2035	DP			
	11083730	Apr 20, 2035	DP U-3118			
	11083797	Apr 20, 2035	DP U-3118			
	11253504	Mar 13, 2034	U-3118			
	11413350	Apr 20, 2035	U-3417			
	11844837	Apr 21, 2036	U-3417			
	9592227	Mar 13, 2034	DP U-3118			
	9694079	Apr 20, 2035	DP U-3118			
	9744163	Mar 13, 2034	DP			
	9801945	Apr 20, 2035	DP U-3118			
	9913909	Mar 13, 2034	U-3118			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>BUPIVACAINE; MELOXICAM - ZYNRELEF KIT</u>						
N 211988 002	10098957	Apr 20, 2035	U-3118		I-933	Jan 23, 2027
	10213510	Apr 20, 2035	DP U-3118			
	10398686	Mar 13, 2034	DP			
	10632199	Apr 20, 2035	DP U-3118			
	10898575	Apr 20, 2035	DP U-3118			
	10980886	Apr 20, 2035	DP			
	11083730	Apr 20, 2035	DP U-3118			
	11083797	Apr 20, 2035	DP U-3118			
	11253504	Mar 13, 2034	U-3118			
	11413350	Apr 20, 2035	U-3417			
	11844837	Apr 21, 2036	U-3417			
	9592227	Mar 13, 2034	DP U-3118			
	9694079	Apr 20, 2035	DP U-3118			
	9744163	Mar 13, 2034	DP			
	9801945	Apr 20, 2035	DP U-3118			
	9913909	Mar 13, 2034	U-3118			
<u>BUPIVACAINE; MELOXICAM - ZYNRELEF KIT</u>						
N 211988 003	10098957	Apr 20, 2035	U-3118		I-933	Jan 23, 2027
	10213510	Apr 20, 2035	DP U-3118			
	10398686	Mar 13, 2034	DP			
	10632199	Apr 20, 2035	DP U-3118			
	10898575	Apr 20, 2035	DP U-3118			
	10980886	Apr 20, 2035	DP			
	11083730	Apr 20, 2035	DP U-3118			
	11083797	Apr 20, 2035	DP U-3118			
	11253504	Mar 13, 2034	U-3118			
	11413350	Apr 20, 2035	U-3417			
	11844837	Apr 21, 2036	U-3417			
	9592227	Mar 13, 2034	DP U-3118			
	9694079	Apr 20, 2035	DP U-3118			
	9744163	Mar 13, 2034	DP			
	9801945	Apr 20, 2035	DP U-3118			
	9913909	Mar 13, 2034	U-3118			
<u>BUPIVACAINE; MELOXICAM - ZYNRELEF KIT</u>						
N 211988 004	10098957	Apr 20, 2035	U-3118		I-933	Jan 23, 2027
	10213510	Apr 20, 2035	DP U-3118			
	10398686	Mar 13, 2034	DP			
	10632199	Apr 20, 2035	DP U-3118			
	10898575	Apr 20, 2035	DP U-3118			
	10980886	Apr 20, 2035	DP			
	11083730	Apr 20, 2035	DP U-3118			
	11083797	Apr 20, 2035	DP U-3118			
	11253504	Mar 13, 2034	U-3118			
	11413350	Apr 20, 2035	U-3417			
	11844837	Apr 21, 2036	U-3417			
	9592227	Mar 13, 2034	DP U-3118			
	9694079	Apr 20, 2035	DP U-3118			
	9744163	Mar 13, 2034	DP			
	9801945	Apr 20, 2035	DP U-3118			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>BUPIVACAINE; MELOXICAM - ZYNRELEF KIT</u>						
N 211988	004 9913909	Mar 13, 2034	U-3118			
<u>BUPRENORPHINE - SUBLOCADE</u>						
N 209819	001 10198218	Jun 06, 2031	U-2489			
	10558394	Jun 25, 2031	DP			
	10592168	Jun 06, 2031	U-2489			
	11000520	Nov 06, 2035	U-3111			
	11839611	Nov 06, 2035	U-3111			
	8921387	Jan 06, 2032	DP U-2173			
	8921387	Jan 06, 2032	DP U-2174			
	8975270	Sep 05, 2031	DP U-2175			
	8975270	Sep 05, 2031	DP U-2206			
	9272044	Jun 06, 2031	U-2176			
	9272044	Jun 06, 2031	U-2177			
	9272044	Jun 06, 2031	U-2178			
	9272044	Jun 06, 2031	U-2209			
	9498432	Jun 06, 2031	DP U-2179			
	9782402	Jun 06, 2031	DP U-2176			
	9782402	Jun 06, 2031	DP U-2180			
	9782402	Jun 06, 2031	DP U-2207			
	9782402	Jun 06, 2031	DP U-2208			
	9827241	Jun 06, 2031	DP U-2174			
	9827241	Jun 06, 2031	DP U-2181			
	9827241	Jun 06, 2031	DP U-2206			
	9827241	Jun 06, 2031	DP U-2210			
	9827241	Jun 06, 2031	DP U-2211			
<u>BUPRENORPHINE - SUBLOCADE</u>						
N 209819	002 10198218	Jun 06, 2031	U-2489			
	10558394	Jun 25, 2031	DP			
	10592168	Jun 06, 2031	U-2489			
	10646484	Jun 22, 2038	U-2489			
	11000520	Nov 06, 2035	U-3111			
	11839611	Nov 06, 2035	U-3111			
	8921387	Jan 06, 2032	DP U-2173			
	8921387	Jan 06, 2032	DP U-2174			
	8975270	Sep 05, 2031	DP U-2175			
	8975270	Sep 05, 2031	DP U-2206			
	9272044	Jun 06, 2031	U-2176			
	9272044	Jun 06, 2031	U-2177			
	9272044	Jun 06, 2031	U-2178			
	9272044	Jun 06, 2031	U-2209			
	9498432	Jun 06, 2031	DP U-2179			
	9782402	Jun 06, 2031	DP U-2176			
	9782402	Jun 06, 2031	DP U-2180			
	9782402	Jun 06, 2031	DP U-2207			
	9782402	Jun 06, 2031	DP U-2208			
	9827241	Jun 06, 2031	DP U-2174			
	9827241	Jun 06, 2031	DP U-2181			
	9827241	Jun 06, 2031	DP U-2206			
	9827241	Jun 06, 2031	DP U-2210			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>BUPRENORPHINE - SUBLOCADE</u>						
N 209819	002 9827241	Jun 06, 2031	DP U-2211			
<u>BUPRENORPHINE - BRIXADI</u>						
N 210136	001 10912772	Jul 26, 2032	U-3617		NP	May 23, 2026
	11110084	Jul 26, 2032	DP U-3616			
	11135215	Jul 26, 2032	DP			
	8236292	Jan 10, 2027	DP U-3619			
	8236755	Jul 31, 2026	DP U-3620			
	8545832	Jun 06, 2025	DP U-3619			
	9937164	Jul 26, 2032	DP U-3618			
<u>BUPRENORPHINE - BRIXADI</u>						
N 210136	002 10912772	Jul 26, 2032	U-3617		NP	May 23, 2026
	11110084	Jul 26, 2032	DP U-3616			
	11135215	Jul 26, 2032	DP			
	8236292	Jan 10, 2027	DP U-3619			
	8236755	Jul 31, 2026	DP U-3620			
	8545832	Jun 06, 2025	DP U-3619			
	9937164	Jul 26, 2032	DP U-3618			
<u>BUPRENORPHINE - BRIXADI</u>						
N 210136	003 10912772	Jul 26, 2032	U-3617		NP	May 23, 2026
	11110084	Jul 26, 2032	DP U-3616			
	11135215	Jul 26, 2032	DP			
	8236292	Jan 10, 2027	DP U-3619			
	8236755	Jul 31, 2026	DP U-3620			
	8545832	Jun 06, 2025	DP U-3619			
	9937164	Jul 26, 2032	DP U-3618			
<u>BUPRENORPHINE - BRIXADI</u>						
N 210136	004 10912772	Jul 26, 2032	U-3617		NP	May 23, 2026
	11110084	Jul 26, 2032	DP U-3616			
	11135215	Jul 26, 2032	DP			
	8236292	Jan 10, 2027	DP U-3619			
	8236755	Jul 31, 2026	DP U-3620			
	8545832	Jun 06, 2025	DP U-3619			
	9937164	Jul 26, 2032	DP U-3618			
<u>BUPRENORPHINE - BRIXADI</u>						
N 210136	005 10912772	Jul 26, 2032	U-3617		NP	May 23, 2026
	11110084	Jul 26, 2032	DP U-3616			
	11135215	Jul 26, 2032	DP			
	8236292	Jan 10, 2027	DP U-3619			
	8236755	Jul 31, 2026	DP U-3620			
	8545832	Jun 06, 2025	DP U-3619			
	9937164	Jul 26, 2032	DP U-3618			
<u>BUPRENORPHINE - BRIXADI</u>						
N 210136	006 10912772	Jul 26, 2032	U-3617		NP	May 23, 2026
	11110084	Jul 26, 2032	DP U-3616			
	11135215	Jul 26, 2032	DP			
	8236292	Jan 10, 2027	DP U-3619			
	8236755	Jul 31, 2026	DP U-3620			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>BUPRENORPHINE - BRIXADI</u>						
N 210136 006	8545832	Jun 06, 2025	DP U-3619			
	9937164	Jul 26, 2032	DP U-3618			
<u>BUPRENORPHINE - BRIXADI</u>						
N 210136 007	10912772	Jul 26, 2032	U-3617		NP	May 23, 2026
	11110084	Jul 26, 2032	DP U-3616			
	11135215	Jul 26, 2032	DP			
	8236292	Jan 10, 2027	DP U-3619			
	8236755	Jul 31, 2026	DP U-3620			
	8545832	Jun 06, 2025	DP U-3619			
	9937164	Jul 26, 2032	DP U-3618			
<u>BUPRENORPHINE HYDROCHLORIDE - BELBUCA</u>						
N 207932 001	8147866	Jul 23, 2027	DP U-1769			
	9655843	Jul 23, 2027	DP U-1556			
	9901539	Dec 21, 2032	U-1556			
<u>BUPRENORPHINE HYDROCHLORIDE - BELBUCA</u>						
N 207932 002	8147866	Jul 23, 2027	DP U-1769			
	9655843	Jul 23, 2027	DP U-1556			
	9901539	Dec 21, 2032	U-1556			
<u>BUPRENORPHINE HYDROCHLORIDE - BELBUCA</u>						
N 207932 003	8147866	Jul 23, 2027	DP U-1769			
	9655843	Jul 23, 2027	DP U-1556			
	9901539	Dec 21, 2032	U-1556			
<u>BUPRENORPHINE HYDROCHLORIDE - BELBUCA</u>						
N 207932 004	8147866	Jul 23, 2027	DP U-1769			
	9655843	Jul 23, 2027	DP U-1556			
	9901539	Dec 21, 2032	U-1556			
<u>BUPRENORPHINE HYDROCHLORIDE - BELBUCA</u>						
N 207932 005	8147866	Jul 23, 2027	DP U-1769			
	9655843	Jul 23, 2027	DP U-1556			
	9901539	Dec 21, 2032	U-1556			
<u>BUPRENORPHINE HYDROCHLORIDE - BELBUCA</u>						
N 207932 006	8147866	Jul 23, 2027	DP U-1769			
	9655843	Jul 23, 2027	DP U-1556			
	9901539	Dec 21, 2032	U-1556			
<u>BUPRENORPHINE HYDROCHLORIDE - BELBUCA</u>						
N 207932 007	8147866	Jul 23, 2027	DP U-1769			
	9655843	Jul 23, 2027	DP U-1556			
	9901539	Dec 21, 2032	U-1556			
<u>BUPRENORPHINE HYDROCHLORIDE; NALOXONE HYDROCHLORIDE - SUBOXONE</u>						
N 022410 001	11135216	Aug 07, 2029	DP U-3111			
	8475832	Mar 26, 2030	DP U-1411			
	9687454	Aug 07, 2029	DP U-1464			
<u>BUPRENORPHINE HYDROCHLORIDE; NALOXONE HYDROCHLORIDE - SUBOXONE</u>						
N 022410 002	11135216	Aug 07, 2029	DP U-3111			
	8475832	Mar 26, 2030	DP U-1411			
	9687454	Aug 07, 2029	DP U-1464			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>BUPRENORPHINE HYDROCHLORIDE; NALOXONE HYDROCHLORIDE - SUBOXONE</u>						
N 022410 002	11135216	Aug 07, 2029	DP U-3111			
	8475832	Mar 26, 2030	DP U-1411			
	9687454	Aug 07, 2029	DP U-1464			
<u>BUPRENORPHINE HYDROCHLORIDE; NALOXONE HYDROCHLORIDE - SUBOXONE</u>						
N 022410 003	11135216	Aug 07, 2029	DP U-3111			
	8475832	Mar 26, 2030	DP U-1411			
	9687454	Aug 07, 2029	DP U-1464			
<u>BUPRENORPHINE HYDROCHLORIDE; NALOXONE HYDROCHLORIDE - SUBOXONE</u>						
N 022410 004	11135216	Aug 07, 2029	DP U-3111			
	8475832	Mar 26, 2030	DP U-1411			
	9687454	Aug 07, 2029	DP U-1464			
<u>BUPRENORPHINE HYDROCHLORIDE; NALOXONE HYDROCHLORIDE - ZUBSOLV</u>						
N 204242 001	10946010	Sep 18, 2032	DP			
	11020388	Sep 18, 2032	DP U-3131			
	11433066	Sep 18, 2032	U-3131			
	8470361	May 22, 2030	DP U-1425			
	8658198	Dec 03, 2027	DP U-1494			
	8940330	Sep 18, 2032	DP			
	9259421	Sep 18, 2032	DP			
	9439900	Sep 18, 2032	DP			
<u>BUPRENORPHINE HYDROCHLORIDE; NALOXONE HYDROCHLORIDE - ZUBSOLV</u>						
N 204242 002	10874661	Sep 18, 2032	DP			
	10946010	Sep 18, 2032	DP			
	11020387	Sep 18, 2032	DP U-3131			
	11020388	Sep 18, 2032	DP U-3131			
	11433066	Sep 18, 2032	U-3131			
	8470361	May 22, 2030	DP U-1425			
	8658198	Dec 03, 2027	DP U-1494			
	8940330	Sep 18, 2032	DP			
	9259421	Sep 18, 2032	DP			
	9439900	Sep 18, 2032	DP			
<u>BUPRENORPHINE HYDROCHLORIDE; NALOXONE HYDROCHLORIDE - ZUBSOLV</u>						
N 204242 003	10946010	Sep 18, 2032	DP			
	11020387	Sep 18, 2032	DP U-3131			
	11020388	Sep 18, 2032	DP U-3131			
	11433066	Sep 18, 2032	U-3131			
	8470361	May 22, 2030	DP U-1425			
	8658198	Dec 03, 2027	DP U-1494			
	8940330	Sep 18, 2032	DP			
	9259421	Sep 18, 2032	DP			
	9439900	Sep 18, 2032	DP			
<u>BUPRENORPHINE HYDROCHLORIDE; NALOXONE HYDROCHLORIDE - ZUBSOLV</u>						
N 204242 004	10946010	Sep 18, 2032	DP			
	11020387	Sep 18, 2032	DP U-3131			
	11020388	Sep 18, 2032	DP U-3131			
	11433066	Sep 18, 2032	U-3131			
	8470361	May 22, 2030	DP U-1425			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>BUPRENORPHINE HYDROCHLORIDE; NALOXONE HYDROCHLORIDE - ZUBSOLV</u>						
N 204242 004	8658198	Dec 03, 2027	DP U-1494			
	8940330	Sep 18, 2032	DP			
	9259421	Sep 18, 2032	DP			
	9439900	Sep 18, 2032	DP			
<u>BUPRENORPHINE HYDROCHLORIDE; NALOXONE HYDROCHLORIDE - ZUBSOLV</u>						
N 204242 005	10946010	Sep 18, 2032	DP			
	11020387	Sep 18, 2032	DP U-3131			
	11020388	Sep 18, 2032	DP U-3131			
	11433066	Sep 18, 2032	U-3131			
	8470361	May 22, 2030	DP U-1425			
	8658198	Dec 03, 2027	DP U-1494			
	8940330	Sep 18, 2032	DP			
	9259421	Sep 18, 2032	DP			
	9439900	Sep 18, 2032	DP			
<u>BUPRENORPHINE HYDROCHLORIDE; NALOXONE HYDROCHLORIDE - ZUBSOLV</u>						
N 204242 006	10946010	Sep 18, 2032	DP			
	11020388	Sep 18, 2032	DP U-3131			
	8470361	May 22, 2030	DP U-1425			
	8658198	Dec 03, 2027	DP U-1494			
	8940330	Sep 18, 2032	DP			
	9259421	Sep 18, 2032	DP			
	9439900	Sep 18, 2032	DP	Y		
<u>BUPRENORPHINE HYDROCHLORIDE; NALOXONE HYDROCHLORIDE - BUNAVAIL</u>						
N 205637 001	8147866	Jul 23, 2027	DP U-1521			
	8703177	Aug 20, 2032	DP			
	9522188	Apr 24, 2035	DP			
	9655843	Jul 23, 2027	DP U-2017			
<u>BUPRENORPHINE HYDROCHLORIDE; NALOXONE HYDROCHLORIDE - BUNAVAIL</u>						
N 205637 002	8147866	Jul 23, 2027	DP U-1521			
	8703177	Aug 20, 2032	DP			
	9522188	Apr 24, 2035	DP			
	9655843	Jul 23, 2027	DP U-2017			
<u>BUPRENORPHINE HYDROCHLORIDE; NALOXONE HYDROCHLORIDE - BUNAVAIL</u>						
N 205637 003	8147866	Jul 23, 2027	DP U-1521			
	8703177	Aug 20, 2032	DP			
	9522188	Apr 24, 2035	DP			
	9655843	Jul 23, 2027	DP U-2017			
<u>BUPROPION HYDROBROMIDE - APLENZIN</u>						
N 022108 001	7241805	Jun 27, 2026	DP			
	7569610	Jun 27, 2026	U-997			
	7572935	Jun 27, 2026	DP			
	7585897	Jun 27, 2026	DP			
	7645802	Jun 27, 2026	DP			
	7649019	Jun 27, 2026	DP			
	7662407	Jun 27, 2026	DP			
	7671094	Jun 27, 2026	DP			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>BUPROPION HYDROBROMIDE - APLENZIN</u>						
N 022108	002	7241805	Jun 27, 2026	DP		
		7569610	Jun 27, 2026	U-997		
		7572935	Jun 27, 2026	DP		
		7585897	Jun 27, 2026	DP		
		7645802	Jun 27, 2026	DP		
		7649019	Jun 27, 2026	DP		
		7662407	Jun 27, 2026	DP		
		7671094	Jun 27, 2026	DP		
<u>BUPROPION HYDROBROMIDE - APLENZIN</u>						
N 022108	003	7241805	Jun 27, 2026	DP		
		7569610	Jun 27, 2026	U-997		
		7572935	Jun 27, 2026	DP		
		7585897	Jun 27, 2026	DP		
		7645802	Jun 27, 2026	DP		
		7649019	Jun 27, 2026	DP		
		7662407	Jun 27, 2026	DP		
		7671094	Jun 27, 2026	DP		
<u>BUPROPION HYDROCHLORIDE - FORFIVO XL</u>						
N 022497	001	7674479	Jun 25, 2027	DP		
<u>BUPROPION HYDROCHLORIDE; DEXTROMETHORPHAN HYDROBROMIDE - AUVELITY</u>						
N 215430	001	10058518	Nov 05, 2034	U-3419	NP	Aug 18, 2025
		10064857	Nov 05, 2034	U-3419		
		10080727	Nov 05, 2034	U-3419		
		10092560	Nov 05, 2034	U-3419		
		10092561	Nov 05, 2034	U-3419		
		10105327	Nov 05, 2034	U-3419		
		10105361	Nov 05, 2034	U-3419		
		10251879	Nov 05, 2034	U-3419		
		10463634	Nov 05, 2034	U-3419		
		10512643	Nov 05, 2034	U-3419		
		10548857	Nov 05, 2034	U-3419		
		10596167	Nov 05, 2034	U-3419		
		10772850	Nov 05, 2034	U-3419		
		10780064	Jan 07, 2040	U-3419		
		10780066	Nov 09, 2034	U-3419		
		10786469	Nov 05, 2034	U-3419		
		10786496	Nov 05, 2034	U-3419		
		10799497	Nov 05, 2034	U-3419		
		10806710	Nov 05, 2034	U-3419		
		10864209	Nov 05, 2034	U-3419		
		10874663	Nov 05, 2034	U-3419		
		10874664	Nov 05, 2034	U-3419		
		10874665	Nov 05, 2034	U-3419		
		10881624	Nov 05, 2034	U-3419		
		10881657	Nov 05, 2034	U-3419		
		10894046	Nov 05, 2034	U-3419		
		10894047	Nov 05, 2034	U-3419		
		10898453	Nov 05, 2034	U-3419		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>BUPROPION HYDROCHLORIDE; DEXTROMETHORPHAN HYDROBROMIDE - AUVELITY</u>						
N 215430 001	10925842	Jan 07, 2040	U-3419			
	10933034	Nov 05, 2034	U-3419			
	10940124	Jan 07, 2040	U-3419			
	10945973	Nov 05, 2034	U-3419			
	10966941	Nov 05, 2034	U-3419			
	10966942	Jan 07, 2040	U-3419			
	10966974	Nov 05, 2034	U-3419			
	11020389	Nov 05, 2034	U-3419			
	11058648	Nov 05, 2034	U-3419			
	11090300	Nov 05, 2034	U-3419			
	11096937	Nov 05, 2034	U-3419			
	11123343	Nov 05, 2034	U-3419			
	11129826	Nov 05, 2034	U-3419			
	11141388	Nov 05, 2034	U-3419			
	11141416	Nov 05, 2034	U-3419			
	11147808	Nov 05, 2034	U-3419			
	11185515	Nov 05, 2034	U-3419			
	11191739	Nov 05, 2034	DP U-3419			
	11197839	Nov 05, 2034	DP U-3419			
	11207281	Nov 05, 2034	U-3419			
	11213521	Nov 05, 2034	U-3419			
	11229640	Nov 05, 2034	U-3419			
	11234946	Nov 05, 2034	U-3419			
	11253491	Nov 05, 2034	U-3419			
	11253492	Nov 05, 2034	U-3419			
	11273133	Nov 05, 2034	U-3419			
	11273134	Nov 05, 2034	U-3419			
	11285118	Nov 05, 2034	U-3419			
	11285146	Nov 05, 2034	U-3419			
	11291638	Nov 05, 2034	U-3419			
	11291665	Nov 05, 2034	U-3419			
	11298351	Nov 05, 2034	U-3419			
	11298352	Nov 05, 2034	U-3419			
	11311534	Nov 05, 2034	U-3419			
	11344544	Nov 05, 2034	U-3419			
	11357744	Nov 05, 2034	U-3419			
	11364233	Nov 05, 2034	U-3419			
	11382874	Nov 05, 2034	U-3419			
	11419867	Nov 05, 2034	U-3419			
	11426370	Nov 05, 2034	U-3419			
	11426401	Nov 05, 2034	U-3419			
	11433067	Nov 05, 2034	DP U-3419			
	11439636	Nov 05, 2034	U-3419			
	11478468	Nov 05, 2034	U-3419			
	11497721	Nov 05, 2034	U-3419			
	11510918	Nov 05, 2034	U-3419			
	11517542	Nov 05, 2034	U-3419			
	11517543	Nov 05, 2034	U-3419			
	11524007	Nov 05, 2034	U-3419			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>BUPROPION HYDROCHLORIDE; DEXTROMETHORPHAN HYDROBROMIDE - AUVELITY</u>						
N 215430 001	11524008	Nov 05, 2034	U-3419			
	11534414	Nov 05, 2034	DP U-3419			
	11541021	Nov 05, 2034	U-3419			
	11541048	Nov 05, 2034	U-3419			
	11596627	Nov 05, 2034	U-3419			
	11617728	Nov 05, 2034	U-3419			
	11617747	Nov 05, 2034	U-3563			
	11717518	Jan 20, 2043	U-3419			
	11730706	Jan 23, 2043	U-3419			
	11752144	Feb 23, 2043	U-3419			
	11779579	Nov 05, 2034	U-3419			
	11839612	Mar 02, 2043	U-3419			
	11844797	Apr 20, 2043	U-3778			
	11883373	Jan 23, 2043	U-3419			
	11896563	Dec 01, 2041	U-3419			
	11925636	Jan 20, 2043	U-3419			
	11969421	Nov 05, 2034	U-3419			
	11986444	Feb 15, 2043	U-3419			
	12036191	Feb 15, 2043	U-3419			
	12042473	Feb 23, 2043	U-3419			
	12109178	Nov 05, 2034	U-3419			
	12146889	Feb 23, 2043	U-3419			
	12156914	Jan 20, 2043	DP			
	8569328	Oct 29, 2033	DP U-3419			
	9168234	Nov 05, 2034	U-3419			
	9198905	Nov 05, 2034	U-3419			
	9205083	Nov 05, 2034	U-3419			
	9238032	Nov 05, 2034	U-3419			
	9278095	Nov 05, 2034	U-3419			
	9314462	Nov 05, 2034	U-3419			
	9370513	Nov 05, 2034	U-3419			
	9375429	Nov 05, 2034	U-3419			
	9408815	Nov 05, 2034	U-3419			
	9421176	Nov 05, 2034	U-3419			
	9457023	Nov 05, 2034	U-3419			
	9457025	Nov 05, 2034	U-3419			
	9474731	Nov 05, 2034	U-3419			
	9486450	Nov 05, 2034	U-3419			
	9700528	Nov 05, 2034	U-3419			
	9700553	Nov 05, 2034	U-3419			
	9707191	Nov 05, 2034	U-3419			
	9763932	Nov 05, 2034	U-3419			
	9861595	Nov 05, 2034	U-3419			
	9867819	Nov 05, 2034	U-3419			
	9968568	Nov 05, 2034	U-3419			
<u>BUPROPION HYDROCHLORIDE; NALTREXONE HYDROCHLORIDE - CONTRAVE</u>						
N 200063 001	10231964	Jul 02, 2034	U-1583			
	10307376	Nov 08, 2027	U-1585			
	10403170	Jun 05, 2033	U-1583			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>BUPROPION HYDROCHLORIDE; NALTREXONE HYDROCHLORIDE - CONTRAVE</u>						
N 200063 001	10828294	Jul 02, 2034	U-1583			
	10835527	Jul 02, 2034	U-1583			
	11033543	Jan 10, 2031	U-1583			
	11139056	Jun 05, 2033	U-1583			
	11324741	May 29, 2029	U-1583			
	11998542	Jul 02, 2034	U-1583			
	12048769	Nov 08, 2027	U-1585			
	7375111	Mar 26, 2025	DP			
	8088786	Feb 03, 2029	DP			
	8318788	Nov 08, 2027	U-1584			
	8722085	Nov 08, 2027	U-1585			
	8916195	Feb 02, 2030	U-1639			
	9107837	Jun 04, 2027	U-1639			
	9125868	Nov 08, 2027	U-1585			
	9248123	Jan 13, 2032	U-1808			
	9633575	Jun 25, 2033	U-1583			
<u>CABAZITAXEL - JEVTANA KIT</u>						
N 201023 001	10583110	Oct 27, 2030	U-2753			
	10716777	Oct 27, 2030	U-2856			
	7241907	Dec 10, 2025	DS			
	7241907*PED	Jun 10, 2026				
	8927592	Oct 27, 2030	U-3200			
	8927592*PED	Apr 27, 2031				
<u>CABOTEGRAVIR - APRETUDE</u>						
N 215499 001	10927129	Apr 28, 2026	DS DP		NCE	Jan 21, 2026
	11224597	Sep 15, 2031	DP			
	12138264	Sep 15, 2031	DP			
	8410103	Apr 28, 2026	DS DP			
<u>CABOTEGRAVIR SODIUM - VOCABRIA</u>						
N 212887 001	10927129	Apr 28, 2026	DS DP		M-273	Jan 31, 2025
	8410103	Apr 28, 2026	DS DP U-3061		NCE	Jan 21, 2026
	8410103	Apr 28, 2026	DS DP U-3348		NPP	Mar 29, 2025
<u>CABOTEGRAVIR; RILPIVIRINE - CABENUVA KIT</u>						
N 212888 001	10927129	Apr 28, 2026	DS DP		D-184	Jan 31, 2025
	11224597	Sep 15, 2031	DP U-3348		NCE	Jan 21, 2026
	11389447	Jun 30, 2027	U-3405		NPP	Mar 29, 2025
	12138264	Sep 15, 2031	DP U-3348			
	7125879	Apr 21, 2025	DS DP U-3348			
	8410103	Apr 28, 2026	DS DP U-3348			
	RE50189	Nov 27, 2031	U-3348			
<u>CABOTEGRAVIR; RILPIVIRINE - CABENUVA KIT</u>						
N 212888 002	10927129	Apr 28, 2026	DS DP		D-184	Jan 31, 2025
	11224597	Sep 15, 2031	DP U-3348		NCE	Jan 21, 2026
	11389447	Jun 30, 2027	U-3405		NPP	Mar 29, 2025
	12138264	Sep 15, 2031	DP U-3348			
	7125879	Apr 21, 2025	DS DP U-3348			
	8410103	Apr 28, 2026	DS DP U-3348			
	RE50189	Nov 27, 2031	U-3348			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>CABOTEGRAVIR; RILPIVIRINE - CABENUVA KIT</u>						
N 212888 002	10927129	Apr 28, 2026	DS DP		D-184	Jan 31, 2025
	11224597	Sep 15, 2031	DP U-3348		NCE	Jan 21, 2026
	11389447	Jun 30, 2027	U-3405		NPP	Mar 29, 2025
	12138264	Sep 15, 2031	DP U-3348			
	7125879	Apr 21, 2025	DS DP U-3348			
	8410103	Apr 28, 2026	DS DP U-3348			
	RE50189	Nov 27, 2031	U-3348			
<u>CABOZANTINIB S-MALATE - COMETRIO</u>						
N 203756 001	11091439	Jan 15, 2030	DS			
	11091440	Jan 15, 2030	DP			
	11098015	Jan 15, 2030	U-1617			
	11298349	Feb 10, 2032	DP			
	12128039	Feb 10, 2032	DP U-1617			
	7579473	Aug 14, 2026	DS DP			
	8877776	Oct 08, 2030	DS DP U-1617			
	9717720	Feb 10, 2032	DP			
<u>CABOZANTINIB S-MALATE - COMETRIO</u>						
N 203756 002	11091439	Jan 15, 2030	DS			
	11091440	Jan 15, 2030	DP			
	11098015	Jan 15, 2030	U-1617			
	11298349	Feb 10, 2032	DP			
	12128039	Feb 10, 2032	DP U-1617			
	7579473	Aug 14, 2026	DS DP			
	8877776	Oct 08, 2030	DS DP U-1617			
	9717720	Feb 10, 2032	DP			
<u>CABOZANTINIB S-MALATE - CABOMETYX</u>						
N 208692 001	10034873	Jul 18, 2031	U-2488		ODE-227	Jan 14, 2026
	10039757	Jul 18, 2031	U-1480		ODE-375	Sep 17, 2028
	11091439	Jan 15, 2030	DS			
	11091440	Jan 15, 2030	DP			
	11098015	Jan 15, 2030	U-1220			
	11098015	Jan 15, 2030	U-1480			
	11098015	Jan 15, 2030	U-2488			
	11098015	Jan 15, 2030	U-3225			
	11298349	Feb 10, 2032	DP			
	12128039	Feb 10, 2032	DP U-1220			
	12128039	Feb 10, 2032	DP U-1480			
	12128039	Feb 10, 2032	DP U-2488			
	12128039	Feb 10, 2032	DP U-3225			
	7579473	Aug 14, 2026	DS DP			
	8877776	Oct 08, 2030	DS DP U-3225			
	9724342	Jul 09, 2033	DP			
<u>CABOZANTINIB S-MALATE - CABOMETYX</u>						
N 208692 002	10034873	Jul 18, 2031	U-2488		ODE-227	Jan 14, 2026
	10039757	Jul 18, 2031	U-1480		ODE-375	Sep 17, 2028
	11091439	Jan 15, 2030	DS			
	11091440	Jan 15, 2030	DP			
	11098015	Jan 15, 2030	U-1220			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>CABOZANTINIB S-MALATE - CABOMETYX</u>						
N 208692 002	11098015	Jan 15, 2030	U-1480			
	11098015	Jan 15, 2030	U-2488			
	11098015	Jan 15, 2030	U-3225			
	11298349	Feb 10, 2032	DP			
	12128039	Feb 10, 2032	DP U-1220			
	12128039	Feb 10, 2032	DP U-1480			
	12128039	Feb 10, 2032	DP U-2488			
	12128039	Feb 10, 2032	DP U-3225			
	7579473	Aug 14, 2026	DS DP			
	8877776	Oct 08, 2030	DS DP U-3225			
	9724342	Jul 09, 2033	DP			
<u>CABOZANTINIB S-MALATE - CABOMETYX</u>						
N 208692 003	10034873	Jul 18, 2031	U-2488		ODE-227	Jan 14, 2026
	10039757	Jul 18, 2031	U-1480		ODE-375	Sep 17, 2028
	11091439	Jan 15, 2030	DS			
	11091440	Jan 15, 2030	DP			
	11098015	Jan 15, 2030	U-1220			
	11098015	Jan 15, 2030	U-1480			
	11098015	Jan 15, 2030	U-2488			
	11098015	Jan 15, 2030	U-3225			
	11298349	Feb 10, 2032	DP			
	12128039	Feb 10, 2032	DP U-1220			
	12128039	Feb 10, 2032	DP U-1480			
	12128039	Feb 10, 2032	DP U-2488			
	12128039	Feb 10, 2032	DP U-3225			
	7579473	Aug 14, 2026	DS DP			
	8877776	Oct 08, 2030	DS DP U-3225			
	9724342	Jul 09, 2033	DP			
<u>CALCIFEDIOL - RAYALDEE</u>						
N 208010 001	10213442	Feb 02, 2027	DP			
	10300078	Mar 14, 2034	DP			
	10357502	Mar 14, 2034	DP			
	11154509	Apr 25, 2028	U-3815			
	11253528	Mar 14, 2034	DP			
	11801253	Sep 07, 2030	DP U-3721			
	11801253	Sep 07, 2030	DP U-3722			
	11801253	Sep 07, 2030	DP U-3723			
	8207149	Apr 25, 2028	U-1871			
	8361488	Jul 19, 2028	DP			
	8426391	Aug 27, 2028	U-1872			
	8778373	Apr 25, 2028	U-1873			
	8906410	Feb 02, 2027	DP			
	9408858	Apr 25, 2028	U-1888			
	9498486	Apr 25, 2028	U-3814			
	9861644	Mar 14, 2034	DP			
	9925147	Apr 25, 2028	DP U-2255			
	9925147	Apr 25, 2028	DP U-2256			
	9925147	Apr 25, 2028	DP U-2257			
	9925147	Apr 25, 2028	DP U-2258			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>CALCIFEDIOL - RAYALDEE</u>						
N 208010	001 9925147	Apr 25, 2028	DP U-2259			
	9943530	Feb 02, 2027	U-2274			
<u>CALCIPOTRIENE - SORILUX</u>						
N 022563	001 8263580	May 07, 2028	DP U-1280			
	8263580	May 07, 2028	DP U-2662			
	8629128	May 26, 2026	DP U-1280			
	8629128	May 26, 2026	DP U-1767			
	8629128	May 26, 2026	DP U-2662			
<u>CALCIUM ACETATE - PHOSLYRA</u>						
N 022581	001 8591938	Feb 23, 2030	DP U-1469			
	8592480	Jul 20, 2027	U-1469			
	9089528	Jul 20, 2027	U-1469			
<u>CALCIUM GLUCONATE - CALCIUM GLUCONATE IN SODIUM CHLORIDE</u>						
N 210906	001 10130646	Jul 25, 2037	DP			
	10342813	Jul 25, 2037	DP			
<u>CALCIUM GLUCONATE - CALCIUM GLUCONATE IN SODIUM CHLORIDE</u>						
N 210906	002 10130646	Jul 25, 2037	DP			
	10342813	Jul 25, 2037	DP			
<u>CALCIUM GLUCONATE - CALCIUM GLUCONATE IN SODIUM CHLORIDE</u>						
N 210906	003 10130646	Jul 25, 2037	DP			
	10342813	Jul 25, 2037	DP			
<u>CALCIUM OXYBATE; MAGNESIUM OXYBATE; POTASSIUM OXYBATE; SODIUM OXYBATE - XYWAV</u>						
N 212690	001 10195168	Jan 11, 2033	DP		ODE-361	Jul 21, 2027
	10213400	Mar 15, 2033	U-2499		ODE-369	Aug 12, 2028
	10213400*PED	Sep 15, 2033			PED	Jan 21, 2028
	10675258	Jan 11, 2033	U-2938			
	10864181	Mar 15, 2033	U-3017			
	10864181*PED	Sep 15, 2033				
	11253494	Mar 15, 2033	U-3323			
	11253494	Mar 15, 2033	U-3324			
	11253494*PED	Sep 15, 2033				
	11426373	Sep 19, 2037	U-3432			
	11554102	Jan 11, 2033	DP			
	11986446	Mar 15, 2033	U-3017			
	11986446	Mar 15, 2033	U-3324			
	12138233	Feb 22, 2041	U-4044			
	8591922	Jan 11, 2033	DP			
	8772306	Mar 15, 2033	U-1532			
	8772306	Mar 15, 2033	U-3198			
	8772306*PED	Sep 15, 2033				
	8901173	Jan 11, 2033	DP			
	9050302	Mar 15, 2033	U-1532			
	9050302*PED	Sep 15, 2033				
	9132107	Jan 11, 2033	DP			
	9486426	Mar 15, 2033	U-1532			
	9486426*PED	Sep 15, 2033				

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>CANAGLIFLOZIN - INVOKANA</u>						
N 204042 001	10617668	May 11, 2031	DP U-2441			
	10617668	May 11, 2031	DP U-2632			
	10617668	May 11, 2031	DP U-2794			
	10617668	May 11, 2031	DP U-2795			
	10617668	May 11, 2031	DP U-2796			
	10617668	May 11, 2031	DP U-2797			
	10617668	May 11, 2031	DP U-2798			
	10617668	May 11, 2031	DP U-2799			
	10617668	May 11, 2031	DP U-493			
	10617668*PED	Nov 11, 2031				
	7943582	Feb 26, 2029	DS DP U-2441			
	7943582	Feb 26, 2029	DS DP U-2632			
	7943582	Feb 26, 2029	DS DP U-493			
	7943582*PED	Aug 26, 2029				
	7943788	Jul 14, 2027	DS DP			
	7943788*PED	Jan 14, 2028				
	8222219	Apr 11, 2025	U-2441			
	8222219	Apr 11, 2025	U-2632			
	8222219	Apr 11, 2025	U-493			
	8222219*PED	Oct 11, 2025				
	8513202	Dec 03, 2027	DS DP U-2441			
	8513202	Dec 03, 2027	DS DP U-2632			
	8513202	Dec 03, 2027	DS DP U-493			
	8513202*PED	Jun 03, 2028				
<u>CANAGLIFLOZIN - INVOKANA</u>						
N 204042 002	10617668	May 11, 2031	DP U-2441			
	10617668	May 11, 2031	DP U-2632			
	10617668	May 11, 2031	DP U-2794			
	10617668	May 11, 2031	DP U-2795			
	10617668	May 11, 2031	DP U-2796			
	10617668	May 11, 2031	DP U-2797			
	10617668	May 11, 2031	DP U-2798			
	10617668	May 11, 2031	DP U-2799			
	10617668	May 11, 2031	DP U-493			
	10617668*PED	Nov 11, 2031				
	7943582	Feb 26, 2029	DS DP U-2441			
	7943582	Feb 26, 2029	DS DP U-2632			
	7943582	Feb 26, 2029	DS DP U-493			
	7943582*PED	Aug 26, 2029				
	7943788	Jul 14, 2027	DS DP			
	7943788*PED	Jan 14, 2028				
	8222219	Apr 11, 2025	U-2441			
	8222219	Apr 11, 2025	U-2632			
	8222219	Apr 11, 2025	U-493			
	8222219*PED	Oct 11, 2025				
	8513202	Dec 03, 2027	DS DP U-2441			
	8513202	Dec 03, 2027	DS DP U-2632			
	8513202	Dec 03, 2027	DS DP U-493			
	8513202*PED	Jun 03, 2028				

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>CANAGLIFLOZIN - INVOKANA</u>						
N 204042 002	10617668	May 11, 2031	DP U-2441			
	10617668	May 11, 2031	DP U-2632			
	10617668	May 11, 2031	DP U-2794			
	10617668	May 11, 2031	DP U-2795			
	10617668	May 11, 2031	DP U-2796			
	10617668	May 11, 2031	DP U-2797			
	10617668	May 11, 2031	DP U-2798			
	10617668	May 11, 2031	DP U-2799			
	10617668	May 11, 2031	DP U-493			
	10617668*PED	Nov 11, 2031				
	7943582	Feb 26, 2029	DS DP U-2441			
	7943582	Feb 26, 2029	DS DP U-2632			
	7943582	Feb 26, 2029	DS DP U-493			
	7943582*PED	Aug 26, 2029				
	7943788	Jul 14, 2027	DS DP			
	7943788*PED	Jan 14, 2028				
	8222219	Apr 11, 2025	U-2441			
	8222219	Apr 11, 2025	U-2632			
	8222219	Apr 11, 2025	U-493			
	8222219*PED	Oct 11, 2025				
	8513202	Dec 03, 2027	DS DP U-2441			
	8513202	Dec 03, 2027	DS DP U-2632			
	8513202	Dec 03, 2027	DS DP U-493			
	8513202*PED	Jun 03, 2028				
<u>CANAGLIFLOZIN; METFORMIN HYDROCHLORIDE - INVOKAMET</u>						
N 204353 001	11576894	Jul 06, 2030	DP			
	7943582	Feb 26, 2029	DS DP U-2441			
	7943582	Feb 26, 2029	DS DP U-2632			
	7943582	Feb 26, 2029	DS DP U-493			
	7943582*PED	Aug 26, 2029				
	7943788	Jul 14, 2027	DS DP			
	7943788*PED	Jan 14, 2028				
	8222219	Apr 11, 2025	U-2441			
	8222219	Apr 11, 2025	U-2632			
	8222219	Apr 11, 2025	U-493			
	8222219*PED	Oct 11, 2025				
	8513202	Dec 03, 2027	DS DP U-2441			
	8513202	Dec 03, 2027	DS DP U-2632			
	8513202	Dec 03, 2027	DS DP U-493			
	8513202*PED	Jun 03, 2028				
<u>CANAGLIFLOZIN; METFORMIN HYDROCHLORIDE - INVOKAMET</u>						
N 204353 002	11576894	Jul 06, 2030	DP			
	7943582	Feb 26, 2029	DS DP U-2441			
	7943582	Feb 26, 2029	DS DP U-2632			
	7943582	Feb 26, 2029	DS DP U-493			
	7943582*PED	Aug 26, 2029				
	7943788	Jul 14, 2027	DS DP			
	7943788*PED	Jan 14, 2028				
	8222219	Apr 11, 2025	U-2441			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>CANAGLIFLOZIN; METFORMIN HYDROCHLORIDE - INVOKAMET</u>						
N 204353 002	8222219	Apr 11, 2025	U-2632			
	8222219	Apr 11, 2025	U-493			
	8222219*PED	Oct 11, 2025				
	8513202	Dec 03, 2027	DS DP U-2441			
	8513202	Dec 03, 2027	DS DP U-2632			
	8513202	Dec 03, 2027	DS DP U-493			
	8513202*PED	Jun 03, 2028				
<u>CANAGLIFLOZIN; METFORMIN HYDROCHLORIDE - INVOKAMET</u>						
N 204353 003	11576894	Jul 06, 2030	DP			
	7943582	Feb 26, 2029	DS DP U-2441			
	7943582	Feb 26, 2029	DS DP U-2632			
	7943582	Feb 26, 2029	DS DP U-493			
	7943582*PED	Aug 26, 2029				
	7943788	Jul 14, 2027	DS DP			
	7943788*PED	Jan 14, 2028				
	8222219	Apr 11, 2025	U-2441			
	8222219	Apr 11, 2025	U-2632			
	8222219	Apr 11, 2025	U-493			
	8222219*PED	Oct 11, 2025				
	8513202	Dec 03, 2027	DS DP U-2441			
	8513202	Dec 03, 2027	DS DP U-2632			
	8513202	Dec 03, 2027	DS DP U-493			
	8513202*PED	Jun 03, 2028				
<u>CANAGLIFLOZIN; METFORMIN HYDROCHLORIDE - INVOKAMET</u>						
N 204353 004	11576894	Jul 06, 2030	DP			
	7943582	Feb 26, 2029	DS DP U-2441			
	7943582	Feb 26, 2029	DS DP U-2632			
	7943582	Feb 26, 2029	DS DP U-493			
	7943582*PED	Aug 26, 2029				
	7943788	Jul 14, 2027	DS DP			
	7943788*PED	Jan 14, 2028				
	8222219	Apr 11, 2025	U-2441			
	8222219	Apr 11, 2025	U-2632			
	8222219	Apr 11, 2025	U-493			
	8222219*PED	Oct 11, 2025				
	8513202	Dec 03, 2027	DS DP U-2441			
	8513202	Dec 03, 2027	DS DP U-2632			
	8513202	Dec 03, 2027	DS DP U-493			
	8513202*PED	Jun 03, 2028				
<u>CANAGLIFLOZIN; METFORMIN HYDROCHLORIDE - INVOKAMET XR</u>						
N 205879 001	7943582	Feb 26, 2029	DS DP U-2441			
	7943582	Feb 26, 2029	DS DP U-2632			
	7943582	Feb 26, 2029	DS DP U-493			
	7943582*PED	Aug 26, 2029				
	7943788	Jul 14, 2027	DS DP			
	7943788*PED	Jan 14, 2028				
	8222219	Apr 11, 2025	U-2441			
	8222219	Apr 11, 2025	U-2632			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>CANAGLIFLOZIN; METFORMIN HYDROCHLORIDE - INVOKAMET XR</u>						
N 205879 001	8222219	Apr 11, 2025	U-493			
	8222219*PED	Oct 11, 2025				
	8513202	Dec 03, 2027	DS DP U-2441			
	8513202	Dec 03, 2027	DS DP U-2632			
	8513202	Dec 03, 2027	DS DP U-493			
	8513202*PED	Jun 03, 2028				
<u>CANAGLIFLOZIN; METFORMIN HYDROCHLORIDE - INVOKAMET XR</u>						
N 205879 002	7943582	Feb 26, 2029	DS DP U-2441			
	7943582	Feb 26, 2029	DS DP U-2632			
	7943582	Feb 26, 2029	DS DP U-493			
	7943582*PED	Aug 26, 2029				
	7943788	Jul 14, 2027	DS DP			
	7943788*PED	Jan 14, 2028				
	8222219	Apr 11, 2025	U-2441			
	8222219	Apr 11, 2025	U-2632			
	8222219	Apr 11, 2025	U-493			
	8222219*PED	Oct 11, 2025				
	8513202	Dec 03, 2027	DS DP U-2441			
	8513202	Dec 03, 2027	DS DP U-2632			
	8513202	Dec 03, 2027	DS DP U-493			
	8513202*PED	Jun 03, 2028				
<u>CANAGLIFLOZIN; METFORMIN HYDROCHLORIDE - INVOKAMET XR</u>						
N 205879 003	7943582	Feb 26, 2029	DS DP U-2441			
	7943582	Feb 26, 2029	DS DP U-2632			
	7943582	Feb 26, 2029	DS DP U-493			
	7943582*PED	Aug 26, 2029				
	7943788	Jul 14, 2027	DS DP			
	7943788*PED	Jan 14, 2028				
	8222219	Apr 11, 2025	U-2441			
	8222219	Apr 11, 2025	U-2632			
	8222219	Apr 11, 2025	U-493			
	8222219*PED	Oct 11, 2025				
	8513202	Dec 03, 2027	DS DP U-2441			
	8513202	Dec 03, 2027	DS DP U-2632			
	8513202	Dec 03, 2027	DS DP U-493			
	8513202*PED	Jun 03, 2028				
<u>CANAGLIFLOZIN; METFORMIN HYDROCHLORIDE - INVOKAMET XR</u>						
N 205879 004	7943582	Feb 26, 2029	DS DP U-2441			
	7943582	Feb 26, 2029	DS DP U-2632			
	7943582	Feb 26, 2029	DS DP U-493			
	7943582*PED	Aug 26, 2029				
	7943788	Jul 14, 2027	DS DP			
	7943788*PED	Jan 14, 2028				
	8222219	Apr 11, 2025	U-2441			
	8222219	Apr 11, 2025	U-2632			
	8222219	Apr 11, 2025	U-493			
	8222219*PED	Oct 11, 2025				
	8513202	Dec 03, 2027	DS DP U-2441			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>CANAGLIFLOZIN; METFORMIN HYDROCHLORIDE - INVOKAMET XR</u>						
N 205879 004	8513202	Dec 03, 2027	DS DP U-2632			
	8513202	Dec 03, 2027	DS DP U-493			
	8513202*PED	Jun 03, 2028				
<u>CANGRELOR - KENGREAL</u>						
N 204958 001	10039780	Jul 10, 2035	U-2260			
	8680052	Mar 09, 2033	U-2979			
	9295687	Jul 10, 2035	DP			
	9427448	Nov 10, 2030	U-1926			
	9439921	Jul 10, 2035	DP			
	9700575	Jul 10, 2035	DP			
	9925265	May 13, 2029	U-2260			
<u>CANNABIDIOL - EPIDIOLEX</u>						
N 210365 001	10092525	Jun 17, 2035	U-2427		M-270	Oct 20, 2026
	10111840	Jun 17, 2035	U-2442		ODE-216	Sep 28, 2025
	10111840	Jun 17, 2035	U-2443		ODE-326	Jul 31, 2027
	10137095	Jun 17, 2035	U-2454		ODE-332	Jul 31, 2027
	10137095	Jun 17, 2035	U-2455			
	10603288	Jun 17, 2035	U-2780			
	10603288	Jun 17, 2035	U-2781			
	10603288	Jun 17, 2035	U-2782			
	10603288	Jun 17, 2035	U-2783			
	10709671	Jun 17, 2035	U-2862			
	10709673	Jun 17, 2035	DP			
	10709674	Jun 17, 2035	U-2780			
	10709674	Jun 17, 2035	U-2781			
	10849860	Jun 17, 2035	U-2427			
	10849860	Jun 17, 2035	U-2454			
	10918608	Oct 13, 2035	U-3071			
	10918608	Oct 13, 2035	U-3072			
	10918608	Oct 13, 2035	U-3073			
	10966939	Jun 17, 2035	DP U-2780			
	10966939	Jun 17, 2035	DP U-2781			
	11065209	Oct 13, 2035	U-3071			
	11096905	Oct 13, 2035	DS DP U-2780			
	11096905	Oct 13, 2035	DS DP U-2781			
	11154516	Jun 17, 2035	U-3235			
	11154516	Jun 17, 2035	U-3236			
	11160795	Mar 01, 2041	U-3233			
	11207292	Apr 26, 2039	DS U-3235			
	11207292	Apr 26, 2039	DS U-3236			
	11207292	Apr 26, 2039	DS U-3277			
	11311498	Jun 17, 2035	U-3375			
	11311498	Jun 17, 2035	U-3376			
	11357741	Jun 17, 2035	U-2862			
	11400055	Oct 13, 2035	U-3071			
	11406623	Mar 01, 2041	U-3233			
	11446258	Jun 17, 2035	U-2780			
	11446258	Jun 17, 2035	U-2781			
	11446258	Jun 17, 2035	U-3071			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>CANNABIDIOL - EPIDIOLEX</u>						
N 210365	001	11633369	Jun 17, 2035	DP U-2780		
		11633369	Jun 17, 2035	DP U-2781		
		11633369	Jun 17, 2035	DP U-3071		
		11701330	Jun 17, 2035	U-2780		
		11701330	Jun 17, 2035	U-2781		
		11766411	Jun 17, 2035	U-2781		
		11865102	Apr 26, 2039	DS DP U-2781		
		11865102	Apr 26, 2039	DS DP U-3236		
		11865102	Apr 26, 2039	DS DP U-3277		
		11963937	Jun 17, 2035	U-2780		
		11963937	Jun 17, 2035	U-2781		
		12064399	Jun 17, 2035	U-3988		
		12064399	Jun 17, 2035	U-3989		
		12102619	Mar 01, 2041	U-3233		
		9949937	Jun 17, 2035	U-2421		
		9956183	Jun 17, 2035	U-2422		
		9956183	Jun 17, 2035	U-2423		
		9956184	Jun 17, 2035	U-2424		
		9956185	Jun 17, 2035	U-2425		
		9956186	Jun 17, 2035	U-2426		
<u>CANTHARIDIN - YCANTH</u>						
N 212905	001	11052064	May 28, 2035	DP U-3663	NCE	Jul 21, 2028
		11052064	May 28, 2035	DP U-3665		
		11147790	Aug 22, 2038	DP U-3663		
		11147790	Aug 22, 2038	DP U-3664		
<u>CAPIVASERTIB - TRUOAP</u>						
N 218197	001	10039766	Apr 16, 2033	U-3762	NCE	Nov 16, 2028
		10059714	Oct 10, 2028	DS DP		
		10654855	Oct 10, 2028	U-3762		
		11760760	Oct 10, 2028	U-3762		
		8101623	Mar 10, 2030	DS DP U-3762		
		8809336	Oct 25, 2025	U-3762		
		9006430	Oct 25, 2025	DP		
		9487525	Apr 16, 2033	DS DP		
<u>CAPIVASERTIB - TRUOAP</u>						
N 218197	002	10039766	Apr 16, 2033	U-3762	NCE	Nov 16, 2028
		10059714	Oct 10, 2028	DS DP		
		10654855	Oct 10, 2028	U-3762		
		11760760	Oct 10, 2028	U-3762		
		8101623	Mar 10, 2030	DS DP U-3762		
		8809336	Oct 25, 2025	U-3762		
		9006430	Oct 25, 2025	DP		
		9487525	Apr 16, 2033	DS DP		
<u>CAPMATINIB HYDROCHLORIDE - TABRECTA</u>						
N 213591	001	10596178	Jul 22, 2035	DS DP	NCE	May 06, 2025
		12084449	Nov 19, 2027	U-2813	ODE-291	May 06, 2027
		7767675	Nov 19, 2032	DS DP		
		8420645	Jun 05, 2031	DS DP		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>CAPMATINIB HYDROCHLORIDE - TABRECTA</u>						
N 213591	001 8461330	Nov 19, 2027	DS DP			
	8901123	May 20, 2029			U-2813	
<u>CAPMATINIB HYDROCHLORIDE - TABRECTA</u>						
N 213591	002 10596178	Jul 22, 2035	DS DP		NCE	May 06, 2025
	12084449	Nov 19, 2027			ODE-291	May 06, 2027
	7767675	Nov 19, 2032	DS DP			
	8420645	Jun 05, 2031	DS DP			
	8461330	Nov 19, 2027	DS DP			
	8901123	May 20, 2029			U-2813	
<u>CAPSAICIN - OUTENZA</u>						
N 022395	001 10034841	Sep 06, 2025	DP			
	8821920	Mar 26, 2030	DP			
	9226903	Dec 15, 2028	DP			
<u>CARBAMAZEPINE - CARNEXIV</u>						
N 206030	001 11529357	Jan 31, 2040	DP			
	7635773	Mar 13, 2029	DP			
	8410077	Mar 13, 2029	DP			
	9493582	Feb 27, 2033	DP			
	9629797	Nov 10, 2028			U-2004	
	9629797	Nov 10, 2028			U-2005	
	9629797	Nov 10, 2028			U-2006	
	9750822	Mar 13, 2029	DP			
	9770407	Nov 10, 2028	DP			
<u>CARBIDOPA; LEVODOPA - RYTARY</u>						
N 203312	001 8377474	Dec 26, 2028	DP U-1645			
	8377474	Dec 26, 2028	DP U-219			
	8454998	Dec 26, 2028	DP U-1645			
	8454998	Dec 26, 2028	DP U-1646			
	8454998	Dec 26, 2028	DP U-1647			
	8454998	Dec 26, 2028	DP U-1649			
	8454998	Dec 26, 2028	DP U-219			
	8557283	Dec 26, 2028	DP U-1645			
	8557283	Dec 26, 2028	DP U-219			
	9089607	Dec 26, 2028	DP U-1645			
	9089607	Dec 26, 2028	DP U-1720			
	9089608	Dec 26, 2028	DP			
	9463246	Dec 26, 2028	DP U-219			
	9533046	Dec 26, 2028	DP U-219			
	9901640	Dec 26, 2028	DP U-219			
<u>CARBIDOPA; LEVODOPA - RYTARY</u>						
N 203312	002 8377474	Dec 26, 2028	DP U-1645			
	8377474	Dec 26, 2028	DP U-219			
	8454998	Dec 26, 2028	DP U-1645			
	8454998	Dec 26, 2028	DP U-1646			
	8454998	Dec 26, 2028	DP U-1647			
	8454998	Dec 26, 2028	DP U-1649			
	8454998	Dec 26, 2028	DP U-219			
	8557283	Dec 26, 2028	DP U-1645			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>CARBIDOPA; LEVODOPA - RYTARY</u>						
N 203312	002	8557283	Dec 26, 2028	DP U-219		
		9089607	Dec 26, 2028	DP U-1645		
		9089607	Dec 26, 2028	DP U-1720		
		9089608	Dec 26, 2028	DP		
		9463246	Dec 26, 2028	DP U-219		
		9533046	Dec 26, 2028	DP U-219		
		9901640	Dec 26, 2028	DP U-219		
<u>CARBIDOPA; LEVODOPA - RYTARY</u>						
N 203312	003	8377474	Dec 26, 2028	DP U-1645		
		8377474	Dec 26, 2028	DP U-219		
		8454998	Dec 26, 2028	DP U-1645		
		8454998	Dec 26, 2028	DP U-1646		
		8454998	Dec 26, 2028	DP U-1647		
		8454998	Dec 26, 2028	DP U-1649		
		8454998	Dec 26, 2028	DP U-219		
		8557283	Dec 26, 2028	DP U-1645		
		8557283	Dec 26, 2028	DP U-219		
		9089607	Dec 26, 2028	DP U-1645		
		9089607	Dec 26, 2028	DP U-1720		
		9089608	Dec 26, 2028	DP		
		9463246	Dec 26, 2028	DP U-219		
		9533046	Dec 26, 2028	DP U-219		
		9901640	Dec 26, 2028	DP U-219		
<u>CARBIDOPA; LEVODOPA - RYTARY</u>						
N 203312	004	8377474	Dec 26, 2028	DP U-1645		
		8377474	Dec 26, 2028	DP U-219		
		8454998	Dec 26, 2028	DP U-1645		
		8454998	Dec 26, 2028	DP U-1646		
		8454998	Dec 26, 2028	DP U-1647		
		8454998	Dec 26, 2028	DP U-1649		
		8454998	Dec 26, 2028	DP U-219		
		8557283	Dec 26, 2028	DP U-1645		
		8557283	Dec 26, 2028	DP U-219		
		9089607	Dec 26, 2028	DP U-1645		
		9089607	Dec 26, 2028	DP U-1720		
		9089608	Dec 26, 2028	DP		
		9463246	Dec 26, 2028	DP U-219		
		9533046	Dec 26, 2028	DP U-219		
		9901640	Dec 26, 2028	DP U-219		
<u>CARBIDOPA; LEVODOPA - DHIVY</u>						
N 214869	001	11033521	Mar 28, 2039	DP U-219		
		11033521	Mar 28, 2039	DP U-3304		
		11033521	Mar 28, 2039	DP U-3305		
		11439613	Mar 28, 2039	U-3557		
		11819485	Mar 28, 2039	U-3557		
<u>CARBIDOPA; LEVODOPA - CREXONT</u>						
N 217186	001	10098845	Oct 07, 2034	DP U-219	NP	Aug 07, 2027
		10292935	Oct 07, 2034	DP U-219		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>CARBIDOPA; LEVODOPA - CREXONT</u>						
N 217186 001	10688058	Oct 07, 2034	DP U-219			
	10973769	Oct 07, 2034	DP U-219			
	10987313	Oct 07, 2034	DP U-219			
	11357733	Oct 07, 2034	DP			
	11622941	Oct 07, 2034	DP			
	11666538	Oct 07, 2034	DP U-219			
	11986449	Dec 21, 2041	U-219			
	12064521	Oct 07, 2034	DP			
	12109185	Dec 21, 2041	U-1649			
	12109185	Dec 21, 2041	U-219			
	12109185	Dec 21, 2041	U-4004			
	12109185	Dec 21, 2041	U-4005			
	12128141	Oct 07, 2034	DP U-219			
	12178918	Oct 07, 2034	DP			
	12178919	Oct 07, 2034	DP U-219			
<u>CARBIDOPA; LEVODOPA - CREXONT</u>						
N 217186 002	10098845	Oct 07, 2034	DP U-219		NP	Aug 07, 2027
	10292935	Oct 07, 2034	DP U-219			
	10688058	Oct 07, 2034	DP U-219			
	10973769	Oct 07, 2034	DP U-219			
	10987313	Oct 07, 2034	DP U-219			
	11357733	Oct 07, 2034	DP			
	11622941	Oct 07, 2034	DP			
	11666538	Oct 07, 2034	DP U-219			
	11986449	Dec 21, 2041	U-219			
	12064521	Oct 07, 2034	DP			
	12109185	Dec 21, 2041	U-1649			
	12109185	Dec 21, 2041	U-219			
	12109185	Dec 21, 2041	U-4004			
	12109185	Dec 21, 2041	U-4005			
	12128141	Oct 07, 2034	DP U-219			
	12178918	Oct 07, 2034	DP			
	12178919	Oct 07, 2034	DP U-219			
<u>CARBIDOPA; LEVODOPA - CREXONT</u>						
N 217186 003	10098845	Oct 07, 2034	DP U-219		NP	Aug 07, 2027
	10292935	Oct 07, 2034	DP U-219			
	10688058	Oct 07, 2034	DP U-219			
	10973769	Oct 07, 2034	DP U-219			
	10987313	Oct 07, 2034	DP U-219			
	11357733	Oct 07, 2034	DP			
	11622941	Oct 07, 2034	DP			
	11666538	Oct 07, 2034	DP U-219			
	11986449	Dec 21, 2041	U-219			
	12064521	Oct 07, 2034	DP			
	12109185	Dec 21, 2041	U-1649			
	12109185	Dec 21, 2041	U-219			
	12109185	Dec 21, 2041	U-4004			
	12109185	Dec 21, 2041	U-4005			
	12128141	Oct 07, 2034	DP U-219			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>CARBIDOPA; LEVODOPA - CREXONT</u>						
N 217186	003 12178918	Oct 07, 2034	DP			
	12178919	Oct 07, 2034	DP U-219			
<u>CARBIDOPA; LEVODOPA - CREXONT</u>						
N 217186	004 10098845	Oct 07, 2034	DP U-219		NP	Aug 07, 2027
	10292935	Oct 07, 2034	DP U-219			
	10688058	Oct 07, 2034	DP U-219			
	10973769	Oct 07, 2034	DP U-219			
	10987313	Oct 07, 2034	DP U-219			
	11357733	Oct 07, 2034	DP			
	11622941	Oct 07, 2034	DP			
	11666538	Oct 07, 2034	DP U-219			
	11986449	Dec 21, 2041	U-219			
	12064521	Oct 07, 2034	DP			
	12109185	Dec 21, 2041	U-1649			
	12109185	Dec 21, 2041	U-219			
	12109185	Dec 21, 2041	U-4004			
	12109185	Dec 21, 2041	U-4005			
	12128141	Oct 07, 2034	DP U-219			
	12178918	Oct 07, 2034	DP			
	12178919	Oct 07, 2034	DP U-219			
<u>CARBINOXAMINE MALEATE - KARBINAL ER</u>						
N 022556	001 8062667	Mar 29, 2029	DP			
	9522191	Jun 15, 2027	DP			
<u>CARFILZOMIB - KYPROLIS</u>						
N 202714	001 7232818	Apr 14, 2025	DS DP			
	7417042	Jul 20, 2026	DS DP			
	7491704	Apr 14, 2025	U-1260			
	7491704	Apr 14, 2025	U-2319			
	7491704	Apr 14, 2025	U-2320			
	7491704	Apr 14, 2025	U-2947			
	7737112	Dec 07, 2027	DP			
	8129346	Apr 14, 2025	U-1260			
	8129346	Apr 14, 2025	U-2319			
	8129346	Apr 14, 2025	U-2320			
	8129346	Apr 14, 2025	U-2947			
	8207125	Apr 14, 2025	DS DP			
	8207126	Apr 14, 2025	DP			
	8207127	Apr 14, 2025	U-1260			
	8207127	Apr 14, 2025	U-2319			
	8207127	Apr 14, 2025	U-2320			
	8207127	Apr 14, 2025	U-2947			
	8207297	Apr 14, 2025	DS DP			
	9493582	Feb 27, 2033	DP			
	RE47954	Oct 21, 2029	U-3449			
<u>CARFILZOMIB - KYPROLIS</u>						
N 202714	002 7232818	Apr 14, 2025	DS DP			
	7417042	Jul 20, 2026	DS DP			
	7491704	Apr 14, 2025	U-1260			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>CARFILZOMIB - KYPROLIS</u>						
N 202714 002	7491704	Apr 14, 2025	U-2319			
	7491704	Apr 14, 2025	U-2320			
	7491704	Apr 14, 2025	U-2947			
	7737112	Dec 07, 2027	DP			
	8129346	Apr 14, 2025	U-1260			
	8129346	Apr 14, 2025	U-2319			
	8129346	Apr 14, 2025	U-2320			
	8129346	Apr 14, 2025	U-2947			
	8207125	Apr 14, 2025	DS DP			
	8207126	Apr 14, 2025	DP			
	8207127	Apr 14, 2025	U-1260			
	8207127	Apr 14, 2025	U-2319			
	8207127	Apr 14, 2025	U-2320			
	8207127	Apr 14, 2025	U-2947			
	8207297	Apr 14, 2025	DS DP			
	9493582	Feb 27, 2033	DP			
	RE47954	Oct 21, 2029	U-3449			
<u>CARFILZOMIB - KYPROLIS</u>						
N 202714 003	7232818	Apr 14, 2025	DS DP			
	7417042	Jul 20, 2026	DS DP			
	7491704	Apr 14, 2025	U-2319			
	7491704	Apr 14, 2025	U-2320			
	7491704	Apr 14, 2025	U-2947			
	7737112	Dec 07, 2027	DP			
	8129346	Apr 14, 2025	U-2319			
	8129346	Apr 14, 2025	U-2320			
	8129346	Apr 14, 2025	U-2947			
	8207125	Apr 14, 2025	DS DP			
	8207126	Apr 14, 2025	DP			
	8207127	Apr 14, 2025	U-2319			
	8207127	Apr 14, 2025	U-2320			
	8207127	Apr 14, 2025	U-2947			
	8207297	Apr 14, 2025	DS DP			
	9493582	Feb 27, 2033	DP			
	RE47954	Oct 21, 2029	U-3449			
<u>CARGLUMIC ACID - CARBAGLU</u>						
N 022562 001					ODE-345	Jan 22, 2028
<u>CARIPRAZINE HYDROCHLORIDE - VRAYLAR</u>						
N 204370 001	7737142	Sep 17, 2029	DS DP U-1750		I-904	Dec 16, 2025
	7737142	Sep 17, 2029	DS DP U-2543			
	7737142	Sep 17, 2029	DS DP U-2544			
	7737142	Sep 17, 2029	DS DP U-2545			
	7737142	Sep 17, 2029	DS DP U-3503			
	7943621	Dec 16, 2028	DS DP			
	RE47350	Jul 16, 2029	U-1750			
	RE47350	Jul 16, 2029	U-2543			
	RE47350	Jul 16, 2029	U-2544			
	RE47350	Jul 16, 2029	U-2545			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>CARIPRAZINE HYDROCHLORIDE - VRAYLAR</u>						
N 204370 001	RE47350	Jul 16, 2029	U-3503			
	RE49110	Jul 16, 2029	U-2543			
	RE49110	Jul 16, 2029	U-2544			
	RE49110	Jul 16, 2029	U-2545			
	RE49110	Jul 16, 2029	U-3503			
	RE49302	Jul 16, 2029	U-2543			
	RE49302	Jul 16, 2029	U-2544			
	RE49302	Jul 16, 2029	U-2545			
	RE49302	Jul 16, 2029	U-3503			
<u>CARIPRAZINE HYDROCHLORIDE - VRAYLAR</u>						
N 204370 002	7737142	Sep 17, 2029	DS DP U-1750		I-904	Dec 16, 2025
	7737142	Sep 17, 2029	DS DP U-2543			
	7737142	Sep 17, 2029	DS DP U-2544			
	7737142	Sep 17, 2029	DS DP U-2545			
	7737142	Sep 17, 2029	DS DP U-3503			
	7943621	Dec 16, 2028	DS DP			
	RE49110	Jul 16, 2029	U-2543			
	RE49110	Jul 16, 2029	U-2544			
	RE49110	Jul 16, 2029	U-2545			
	RE49110	Jul 16, 2029	U-3503			
	RE49302	Jul 16, 2029	U-2543			
	RE49302	Jul 16, 2029	U-2544			
	RE49302	Jul 16, 2029	U-2545			
	RE49302	Jul 16, 2029	U-3503			
<u>CARIPRAZINE HYDROCHLORIDE - VRAYLAR</u>						
N 204370 003	7737142	Sep 17, 2029	DS DP U-1750		I-904	Dec 16, 2025
	7737142	Sep 17, 2029	DS DP U-2543			
	7737142	Sep 17, 2029	DS DP U-2544			
	7943621	Dec 16, 2028	DS DP			
	RE49110	Jul 16, 2029	U-2543			
	RE49110	Jul 16, 2029	U-2544			
	RE49110	Jul 16, 2029	U-2545			
	RE49302	Jul 16, 2029	U-2543			
	RE49302	Jul 16, 2029	U-2544			
	RE49302	Jul 16, 2029	U-2545			
	RE49302	Jul 16, 2029	U-3503			
<u>CARIPRAZINE HYDROCHLORIDE - VRAYLAR</u>						
N 204370 004	7737142	Sep 17, 2029	DS DP U-1750		I-904	Dec 16, 2025
	7737142	Sep 17, 2029	DS DP U-2543			
	7737142	Sep 17, 2029	DS DP U-2544			
	7943621	Dec 16, 2028	DS DP			
	RE49110	Jul 16, 2029	U-2543			
	RE49110	Jul 16, 2029	U-2544			
	RE49110	Jul 16, 2029	U-2545			
	RE49302	Jul 16, 2029	U-2543			
	RE49302	Jul 16, 2029	U-2544			
	RE49302	Jul 16, 2029	U-2545			
	RE49302	Jul 16, 2029	U-3503			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>CARVEDILOL PHOSPHATE - COREG CR</u>						
N 022012 001	8101209	Sep 11, 2025	DP			
<u>CARVEDILOL PHOSPHATE - COREG CR</u>						
N 022012 002	8101209	Sep 11, 2025	DP			
<u>CARVEDILOL PHOSPHATE - COREG CR</u>						
N 022012 003	8101209	Sep 11, 2025	DP			
<u>CARVEDILOL PHOSPHATE - COREG CR</u>						
N 022012 004	8101209	Sep 11, 2025	DP			
<u>CASIMERSEN - AMONDYS 45</u>						
N 213026 001	10287586	Nov 12, 2030	DS DP		NCE	Feb 25, 2026
	10781450	Nov 12, 2030		U-3089	ODE-347	Feb 25, 2028
	9228187	Nov 12, 2030	DS DP			
	9447415	Jun 28, 2025	DS DP			
	9758783	Nov 12, 2030		U-3088		
	9758783	Nov 12, 2030		U-3089		
	RE48960	Jun 28, 2025	DS DP	U-3087		
	RE48960	Jun 28, 2025	DS DP	U-3088		
<u>CASPOFUNGIN ACETATE - CASPOFUNGIN ACETATE</u>						
N 206110 001	9636407	Dec 21, 2032	DP			
<u>CASPOFUNGIN ACETATE - CASPOFUNGIN ACETATE</u>						
N 206110 002	9636407	Dec 21, 2032	DP			
<u>CEDAZURIDINE; DECITABINE - INQOVI</u>						
N 212576 001	11963971	Feb 24, 2041	DP		NCE	Jul 07, 2025
	8268800	Aug 22, 2030	DS	U-2864	ODE-316	Jul 07, 2027
	8268800	Aug 22, 2030	DS	U-2865		
	8268800	Aug 22, 2030	DS	U-2866		
	8268800	Aug 22, 2030	DS	U-2867		
	8618075	Oct 16, 2028		U-2864		
	8618075	Oct 16, 2028		U-2867		
	9567363	Oct 16, 2028	DS			
<u>CEFEPIME HYDROCHLORIDE; ENMETAZOBACTAM - EXBLIFEP</u>						
N 216165 001	11124526	Nov 07, 2034		U-3852	NCE	Feb 22, 2029
	7687488	Dec 03, 2027	DS DP	U-3851	GAIN	Feb 22, 2034
<u>CEFIDEROCOL SULFATE TOSYLATE - FETROJA</u>						
N 209445 001	10004750	Sep 03, 2035	DS DP		NCE	Nov 14, 2024
	9238657	Nov 14, 2033	DS DP	U-282	GAIN	Nov 14, 2029
	9238657	Nov 14, 2033	DS DP	U-3470		
	9238657	Nov 14, 2033	DS DP	U-3471		
	9949982	Sep 03, 2035	DP			
<u>CEFIXIME - SUPRAX</u>						
N 202091 001	9233112	Dec 14, 2028	DP	U-1676		
<u>CEFTAROLINE FOSAMIL - TEFLARO</u>						
N 200327 001	8247400	Feb 10, 2031	DP	U-282		
	9629861	Sep 21, 2030	DP			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>CEFTAROLINE FOSAMIL - TEFLARO</u>						
N 200327	002 8247400	Feb 10, 2031	DP U-282			
	9629861	Sep 21, 2030	DP			
<u>CEFTOBIPROLE MEDOCARIL SODIUM - ZEVTERA</u>						
N 218275	001				NCE	Apr 03, 2029
					GAIN	Apr 03, 2034
<u>CEFTOLOZANE SULFATE; TAZOBACTAM SODIUM - ZERBAXA</u>						
N 206829	001 10028963	Sep 07, 2032	U-2565		NPP	Apr 21, 2025
	10028963	Sep 07, 2032	U-2566			
	10125149	Aug 14, 2035	DP			
	10376496	Sep 09, 2034	U-2610			
	10376496	Sep 09, 2034	U-2611			
	10420841	Mar 14, 2034	U-1672			
	10420841	Mar 14, 2034	U-2631			
	10420841	Mar 14, 2034	U-3360			
	10420841	Mar 14, 2034	U-3361			
	10933053	Sep 09, 2034	U-3090			
	10933053	Sep 09, 2034	U-3091			
	11278622	Mar 14, 2034	U-3335			
	11278622	Mar 14, 2034	U-3336			
	7129232	May 15, 2028	DS DP U-1676			
	7129232	May 15, 2028	DS DP U-3360			
	7129232	May 15, 2028	DS DP U-3361			
	7129232	May 15, 2028	DS DP U-36			
	8476425	Sep 27, 2032	DS			
	8685957	Sep 27, 2032	DS U-36			
	8906898	May 28, 2034	DS DP			
	8968753	Mar 14, 2034	U-1672			
	8968753	Mar 14, 2034	U-1673			
	8968753	Mar 14, 2034	U-3360			
	8968753	Mar 14, 2034	U-3361			
	9320740	Mar 14, 2034	DP			
	9724353	Sep 07, 2032	U-2565			
	9724353	Sep 07, 2032	U-2566			
	9872906	Mar 14, 2034	DP			
<u>CELECOXIB - ELYXYB</u>						
N 212157	001 10376527	May 27, 2036	DP U-2718			
	10722456	May 27, 2036	DP U-2718			
	10799517	May 27, 2036	DP U-2718			
	9572819	May 27, 2036	DP U-2718			
	9795620	May 27, 2036	DP U-2718			
	9949990	May 27, 2036	DP U-2718			
<u>CELECOXIB; TRAMADOL HYDROCHLORIDE - SEGLENTIS</u>						
N 213426	001 10238668	Apr 19, 2030	DS DP U-3244			
	10245276	Apr 19, 2030	DS DP			
	10548909	Apr 19, 2030	U-3244			
	11478488	Apr 19, 2030	U-3244			
	8598152	Apr 19, 2030	DS DP			
	8846744	Jun 03, 2031	DP			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>CELECOXIB; TRAMADOL HYDROCHLORIDE - SEGLENTIS</u>						
N 213426	001 9012440	Apr 19, 2030	DS DP			
<u>CENOBAMATE - XCOPRI</u>						
N 212839	001 11654133	Jun 16, 2039		U-3610	NCE	Mar 10, 2025
	7598279	Oct 30, 2032	DS			
<u>CENOBAMATE - XCOPRI</u>						
N 212839	002 11654133	Jun 16, 2039		U-3610	NCE	Mar 10, 2025
	7598279	Oct 30, 2032	DS			
<u>CENOBAMATE - XCOPRI</u>						
N 212839	003 11654133	Jun 16, 2039		U-3610	NCE	Mar 10, 2025
	7598279	Oct 30, 2032	DS			
<u>CENOBAMATE - XCOPRI</u>						
N 212839	004 11654133	Jun 16, 2039		U-3610	NCE	Mar 10, 2025
	7598279	Oct 30, 2032	DS			
<u>CENOBAMATE - XCOPRI</u>						
N 212839	005 11654133	Jun 16, 2039		U-3610	NCE	Mar 10, 2025
	7598279	Oct 30, 2032	DS			
<u>CENOBAMATE - XCOPRI</u>						
N 212839	006 11654133	Jun 16, 2039		U-3610	NCE	Mar 10, 2025
	7598279	Oct 30, 2032	DS			
<u>CEPHALEXIN - CEPHALEXIN</u>						
A 218817	001				CGT	Apr 09, 2025
<u>CEPHALEXIN - CEPHALEXIN</u>						
A 218817	002				CGT	Apr 09, 2025
<u>CERITINIB - ZYKADIA</u>						
N 205755	001 7893074	Apr 25, 2026	DS DP			
	7964592	Apr 29, 2028	DS DP			
	8039479	Jun 29, 2030	DS DP			
	8377921	Nov 20, 2027		U-1179		
	8399450	Nov 20, 2027	DS DP			
	8703787	Feb 02, 2032		U-1179		
	9309229	Jan 18, 2032	DS DP			
<u>CERITINIB - ZYKADIA</u>						
N 211225	001 7893074	Apr 25, 2026	DS DP			
	7964592	Apr 29, 2028	DS DP			
	8039479	Jun 29, 2030	DS DP			
	8377921	Nov 20, 2027		U-1179		
	8399450	Nov 20, 2027	DS DP			
	8703787	Feb 02, 2032		U-1179		
	9309229	Jan 18, 2032	DS DP			
<u>CETIRIZINE HYDROCHLORIDE - ZERVIAE</u>						
N 208694	001 8829005	Mar 15, 2030		U-1680		
	8829005*PED	Sep 15, 2030				
	9254286	Jul 09, 2032	DP			
	9254286*PED	Jan 09, 2033				
	9750684	Mar 15, 2030	DP			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>CETIRIZINE HYDROCHLORIDE - ZERVIAE</u>						
N 208694	001 9993471	Mar 15, 2030	U-1680			
<u>CETIRIZINE HYDROCHLORIDE - OUZYTTIR</u>						
N 211415	001 8263581	Feb 28, 2030	U-2635			
	8314083	Feb 28, 2030	U-2634			
	8513259	Feb 11, 2030	U-2636			
	9119771	Feb 11, 2030	U-2635			
	9180090	Feb 11, 2030	U-2635			
<u>CHLORHEXIDINE GLUCONATE - CHLORHEXIDINE GLUCONATE</u>						
N 021669	001 7427574	Apr 25, 2026	DP			
	7717889	Feb 27, 2025	DP U-1022			
	7935093	Oct 02, 2027	DP U-1022			
<u>CHLORHEXIDINE GLUCONATE; ISOPROPYL ALCOHOL - CHLORAPREP WITH TINT</u>						
N 020832	002 7422388	Apr 25, 2027	DP U-1397			
<u>CHLORHEXIDINE GLUCONATE; ISOPROPYL ALCOHOL - CHLORAPREP WITH TINT</u>						
N 020832	005 7422388	Apr 25, 2027	DP U-1397			
<u>CHLORHEXIDINE GLUCONATE; ISOPROPYL ALCOHOL - CHLORAPREP WITH TINT</u>						
N 020832	007 7422388	Apr 25, 2027	DP U-1397			
<u>CHLORHEXIDINE GLUCONATE; ISOPROPYL ALCOHOL - SOLUPREP S</u>						
N 208288	001				M-292	Jan 20, 2026
<u>CHLOROPROCAINE HYDROCHLORIDE - CLOROTEKAL</u>						
N 208791	001 8969412	Sep 05, 2026	DP U-2609			
	9504666	Dec 11, 2033	DP			
<u>CHLOROPROCAINE HYDROCHLORIDE - IHEEZO</u>						
N 216227	001 10792271	Sep 15, 2038	DP U-3457		NP	Sep 27, 2025
<u>CHLORPHENIRAMINE MALEATE; CODEINE PHOSPHATE - TUXARIN ER</u>						
N 206323	001 9066942	Jan 03, 2032	U-1716			
	9107921	Jan 03, 2032	DP			
<u>CHLORPHENIRAMINE MALEATE; IBUPROFEN; PSEUDOEPHEDRINE HYDROCHLORIDE - ADVIL ALLERGY SINUS</u>						
N 021441	001 7863287	Feb 28, 2027	DP			
<u>CHLORPHENIRAMINE POLISTIREX; CODEINE POLISTIREX - TUZISTRA XR</u>						
N 207768	001 8062667	Mar 29, 2029	DP			
	8790700	Mar 15, 2027	DP			
<u>CHOLINE FENOFIBRATE - TRILIPIX</u>						
N 022224	001 7259186	Jan 07, 2025	DS			
<u>CHOLINE FENOFIBRATE - TRILIPIX</u>						
N 022224	002 7259186	Jan 07, 2025	DS			
<u>CICLESONIDE - ALVESCO</u>						
N 021658	002 8371292	Feb 01, 2028	U-1355			
<u>CICLESONIDE - ALVESCO</u>						
N 021658	003 8371292	Feb 01, 2028	U-1355			
<u>CICLESONIDE - OMNARIS</u>						
N 022004	001 8371292	Feb 01, 2028	U-1356			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>CICLESONIDE - ZETONNA</u>						
N 202129	001 8371292	Feb 01, 2028	U-1357			
<u>CILASTATIN SODIUM; IMIPENEM; RELEBACTAM - RECARBRIO</u>						
N 212819	001 8487093	Mar 21, 2033	DS DP U-2586		NCE	Jul 16, 2024
	8487093	Mar 21, 2033	DS DP U-2587		GAIN	Jul 16, 2029
	8487093	Mar 21, 2033	DS DP U-2840			
<u>CINACALCET HYDROCHLORIDE - SENSIPAR</u>						
N 021688	001 7829595	Sep 22, 2026	DP U-1098			
	9375405	Sep 22, 2026	DP			
<u>CINACALCET HYDROCHLORIDE - SENSIPAR</u>						
N 021688	002 7829595	Sep 22, 2026	DP U-1098			
	9375405	Sep 22, 2026	DP			
<u>CINACALCET HYDROCHLORIDE - SENSIPAR</u>						
N 021688	003 7829595	Sep 22, 2026	DP U-1098			
	9375405	Sep 22, 2026	DP			
<u>CIPROFLOXACIN - OTIPRIO</u>						
N 207986	001 11040004	Nov 12, 2037	U-2252			
	11246863	Nov 27, 2038	DP			
	11369566	Apr 21, 2029	DP			
	8318817	Apr 27, 2030	U-1792			
	9205048	Apr 21, 2029	U-1793			
	9220796	Jul 01, 2035	DP			
	9233068	Dec 11, 2029	DP			
	9603796	Apr 21, 2029	DS DP U-2252			
<u>CIPROFLOXACIN HYDROCHLORIDE - CIPROFLOXACIN HYDROCHLORIDE</u>						
A 217887	001				CGT	Jun 11, 2025
<u>CIPROFLOXACIN HYDROCHLORIDE; FLUOCINOLONE ACETONIDE - OTOVEL</u>						
N 208251	001 8932610	Mar 24, 2030	DP U-1578			
<u>CIPROFLOXACIN; DEXAMETHASONE - CIPRODEX</u>						
N 021537	001 8846650	Jun 04, 2025	DP U-1578			
<u>CITRIC ACID; LACTIC ACID; POTASSIUM BITARTRATE - PHEXXI</u>						
N 208352	001 10568855	Mar 15, 2033	U-1			
	11337989	Mar 15, 2033	U-1			
	11439610	Mar 15, 2033	DS DP			
	11992472	Mar 15, 2033	U-1			
<u>CITRIC ACID; MAGNESIUM OXIDE; SODIUM PICOSULFATE - PREPOPIK</u>						
N 202535	001 8450338	Oct 10, 2028	DP			
	8481083	Oct 10, 2028	DP			
<u>CITRIC ACID; MAGNESIUM OXIDE; SODIUM PICOSULFATE - CLENPIO</u>						
N 209589	001 10624879	Jun 23, 2034	DP			
	11191753	Jun 23, 2034	U-3261			
	9827231	Jun 26, 2034	DP U-2162			
<u>CLADRIBINE - MAVENCLAD</u>						
N 022561	001 10849919	Nov 23, 2038	U-3411			
	7713947	Oct 16, 2026	U-2520			
	8377903	May 31, 2026	U-2522			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>CLADRIBINE - MAVENCLAD</u>						
N 022561	001 10849919	Nov 23, 2038	U-3411			
	7713947	Oct 16, 2026	U-2520			
	8377903	May 31, 2026	U-2522			
<u>CLASCOTERONE - WINLEVI</u>						
N 213433	001 10159682	Aug 14, 2028	U-2942		NCE	Aug 26, 2025
	11207332	Nov 20, 2028	DP U-3280			
	11938141	Jul 24, 2028	DP			
	8143240	Jan 12, 2026	U-2942			
	8785427	Jul 25, 2030	DP			
	8865690	Jul 24, 2025	U-2942			
	9211295	May 31, 2025	DP			
	9433628	Feb 28, 2029	DP			
	9486458	Jul 24, 2028	U-2942			
<u>CLEVIDIPINE - CLEVIPREX</u>						
N 022156	001 10010537	Oct 10, 2031	DP			
	11103490	Oct 10, 2031	DP			
	8658676	Oct 10, 2031	DP			
<u>CLEVIDIPINE - CLEVIPREX</u>						
N 022156	002 10010537	Oct 10, 2031	DP			
	11103490	Oct 10, 2031	DP			
	8658676	Oct 10, 2031	DP			
<u>CLEVIDIPINE - CLEVIPREX</u>						
N 022156	003 10010537	Oct 10, 2031	DP			
	11103490	Oct 10, 2031	DP			
	8658676	Oct 10, 2031	DP			
<u>CLINDAMYCIN PHOSPHATE - CLINDESSE</u>						
N 050793	001 9789057	Dec 02, 2026	DP U-137			
<u>CLINDAMYCIN PHOSPHATE - XACIATO</u>						
N 215650	001 11129896	Sep 22, 2036	U-3293		NP GAIN	Dec 07, 2024 Dec 07, 2029
<u>CLOBAZAM - SYMPAZAN</u>						
N 210833	001 11541002	Jan 31, 2040	DP U-724			
<u>CLOBAZAM - SYMPAZAN</u>						
N 210833	002 11541002	Jan 31, 2040	DP U-724			
<u>CLOBAZAM - SYMPAZAN</u>						
N 210833	003 11541002	Jan 31, 2040	DP U-724			
<u>CLOBETASOL PROPIONATE - OLUX E</u>						
N 022013	001 8460641	Nov 05, 2028	DP U-1410			
	8962000	Aug 31, 2025	DP U-1410			
<u>CLOBETASOL PROPIONATE - IMPOYZ</u>						
N 209483	001 10064875	Aug 31, 2030	DP U-1408			
	10064875	Aug 31, 2030	DP U-1858			
	10064875	Aug 31, 2030	DP U-193			
	10064875	Aug 31, 2030	DP U-742			
	10064875	Aug 31, 2030	DP U-88			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>CLOBETASOL PROPIONATE - IMPOYZ</u>						
N 209483	001 10588914	Aug 31, 2030	DP U-2771			
	9855334	Mar 11, 2035	DP			
	9956231	Aug 31, 2030	DP U-1408			
	9956231	Aug 31, 2030	DP U-1761			
	9956231	Aug 31, 2030	DP U-1858			
	9956231	Aug 31, 2030	DP U-193			
	9956231	Aug 31, 2030	DP U-742			
	9956231	Aug 31, 2030	DP U-88			
<u>CLOBETASOL PROPIONATE - CLOBETASOL PROPIONATE</u>						
N 218158	001 10588913	May 09, 2036	DP		NP	Mar 04, 2027
	11376262	May 09, 2036	DP U-19			
<u>CLOMIPHENE CITRATE - CLOMIPHENE CITRATE</u>						
A 216739	001				CGT	May 18, 2025
<u>CLONIDINE - NEXICLON XR</u>						
N 022500	001 8337890	Apr 17, 2027	DP			
	8623409	Sep 08, 2031	DP			
<u>CLONIDINE - NEXICLON XR</u>						
N 022500	002 8337890	Apr 17, 2027	DP			
	8623409	Sep 08, 2031	DP			
<u>CLONIDINE HYDROCHLORIDE - ONYDA XR</u>						
N 217645	001 11918689	Jul 28, 2041	DP U-3944			
	8062667	Mar 29, 2029	DP			
<u>CLOZAPINE - VERSACLOZ</u>						
N 203479	001 8057811	May 01, 2028	DP			
<u>COBICISTAT - TYBOST</u>						
N 203094	001 10039718	Oct 06, 2032	DP		ODE-260	Aug 22, 2026
	10039718*PED	Apr 06, 2033				
	8148374	Sep 03, 2029	DS DP U-1279			
	8148374*PED	Mar 03, 2030				
<u>COBICISTAT; DARUNAVIR - PREZCOBIX</u>						
N 205395	001 10039718	Oct 06, 2032	DP			
	7700645	Dec 26, 2026	DS DP			
	7700645*PED	Jun 26, 2027				
	8148374	Sep 03, 2029	DS DP U-1279			
	8148374	Sep 03, 2029	DS DP U-2939			
<u>COBICISTAT; DARUNAVIR; EMTRICITABINE; TENOFOVIR ALAFENAMIDE FUMARATE - SYMTUZA</u>						
N 210455	001 10039718	Oct 06, 2032	DP			
	10786518	Jul 19, 2038	U-2978			
	7390791	Apr 17, 2025	DS DP			
	7700645	Dec 26, 2026	DS DP			
	8148374	Sep 03, 2029	DS DP U-2353			
	8148374	Sep 03, 2029	DS DP U-2364			
	8148374	Sep 03, 2029	DS DP U-2365			
	8148374	Sep 03, 2029	DS DP U-2766			
	8148374	Sep 03, 2029	DS DP U-2767			
	8148374	Sep 03, 2029	DS DP U-2768			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>COBICISTAT; DARUNAVIR; EMTRICITABINE; TENOFOVIR ALAFENAMIDE FUMARATE - SYMTUZA</u>						
N 210455	001	8754065	Aug 15, 2032	DS DP U-2352		
		8754065	Aug 15, 2032	DS DP U-2765		
		9296769	Aug 15, 2032	DS DP U-2352		
		9296769	Aug 15, 2032	DS DP U-2765		
<u>COBICISTAT; ELVITEGRAVIR; EMTRICITABINE; TENOFOVIR ALAFENAMIDE FUMARATE - GENVOYA</u>						
N 207561	001	10039718	Oct 06, 2032	DP		
		10039718*PED	Apr 06, 2033			
		7176220	Aug 27, 2026	DS DP U-257		
		7176220*PED	Feb 27, 2027			
		7390791	Apr 17, 2025	DS DP		
		7390791*PED	Oct 17, 2025			
		7635704	Oct 26, 2026	DS DP U-257		
		7635704*PED	Apr 26, 2027			
		8148374	Sep 03, 2029	DS DP U-1279		
		8148374*PED	Mar 03, 2030			
		8633219	Apr 30, 2030	DP U-257		
		8633219*PED	Oct 30, 2030			
		8754065	Aug 15, 2032	DS DP U-257		
		8754065*PED	Feb 15, 2033			
		8981103	Oct 26, 2026	DS DP		
		8981103*PED	Apr 26, 2027			
		9296769	Aug 15, 2032	DS DP U-257		
		9296769*PED	Feb 15, 2033			
		9891239	Sep 03, 2029	DP U-257		
		9891239*PED	Mar 03, 2030			
<u>COBICISTAT; ELVITEGRAVIR; EMTRICITABINE; TENOFOVIR DISOPROXIL FUMARATE - STRIBILD</u>						
N 203100	001	10039718	Oct 06, 2032	DP		
		10039718*PED	Apr 06, 2033			
		7176220	Aug 27, 2026	DS DP U-257		
		7176220*PED	Feb 27, 2027			
		7635704	Oct 26, 2026	DS DP U-257		
		7635704*PED	Apr 26, 2027			
		8148374	Sep 03, 2029	DS DP U-1279		
		8633219	Apr 30, 2030	DP U-257		
		8633219*PED	Oct 30, 2030			
		8981103	Oct 26, 2026	DS DP		
		8981103*PED	Apr 26, 2027			
		9891239	Sep 03, 2029	DP U-257		
		9891239*PED	Mar 03, 2030			
<u>COBIMETINIB FUMARATE - COTELLIC</u>						
N 206192	001	10478400	Jun 29, 2036	DS DP U-1776	I-902	Oct 28, 2025
		10478400*PED	Dec 29, 2036		M-278	Jul 28, 2025
		10590102	Jun 30, 2036	DS DP U-1776	ODE-416	Oct 28, 2029
		10590102*PED	Dec 30, 2036		PED	Jan 28, 2026
		11087354	Jun 22, 2034	U-1776		
		11087354*PED	Dec 22, 2034			
		11254649	Jun 30, 2036	DS DP U-1776		
		11254649*PED	Dec 30, 2036			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>COBIMETINIB FUMARATE - COTELLIC</u>						
N 206192	001 11597699	Oct 05, 2026				U-3554
	7803839	Nov 10, 2029	DS DP			
	7803839*PED	May 10, 2030				
	8362002	Oct 05, 2026				U-1776
	8362002*PED	Apr 05, 2027				
<u>COCAINE HYDROCHLORIDE - NUMBRINO</u>						
N 209575	001 10016407	Feb 07, 2037				U-2225
	10016407	Feb 07, 2037				U-2226
	10016407	Feb 07, 2037				U-2329
	10149843	Feb 07, 2037				U-2478
	10149843	Feb 07, 2037				U-2479
	10231961	Feb 07, 2037	DP			
	10413505	Feb 07, 2037				U-3014
	10420760	Feb 07, 2037				U-2478
	10857095	Feb 07, 2037				U-3014
	10894012	Feb 07, 2037				U-3014
	10933060	Feb 07, 2037				U-2478
	10933060	Feb 07, 2037				U-3014
	10973811	Feb 07, 2037				U-3680
	11040032	Feb 07, 2037	DP			
	9867815	Feb 07, 2037				U-2225
	9867815	Feb 07, 2037				U-2226
	9867815	Feb 07, 2037				U-2227
	9867815	Feb 07, 2037				U-2329
<u>COCAINE HYDROCHLORIDE - GOPRELTO</u>						
N 209963	001 10016407	Feb 07, 2037				U-2329
	10149843	Feb 07, 2037				U-2478
	10149843	Feb 07, 2037				U-2479
	10231961	Feb 07, 2037	DP			
	10413505	Feb 07, 2037				U-2479
	10420760	Feb 07, 2037				U-2478
	10857095	Feb 07, 2037				U-3014
	10894012	Feb 07, 2037				U-3014
	10933060	Feb 07, 2037				U-3014
	10973811	Feb 07, 2037				U-2226
	10987347	Feb 07, 2037				U-2225
	11040032	Feb 07, 2037	DP			
	9867815	Feb 07, 2037				U-2225
	9867815	Feb 07, 2037				U-2226
	9867815	Feb 07, 2037				U-2227
<u>COLCHICINE - COLCRYS</u>						
N 022352	001 7601758	Feb 10, 2029				U-1007
	7619004	Dec 03, 2028				U-1020
	7820681	Feb 17, 2029				U-1020
	7906519	Feb 17, 2029				U-1116
	7915269	Feb 17, 2029				U-1007
	7935731	Dec 03, 2028				U-1116
	7964647	Oct 06, 2028				U-1007

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>COLCHICINE - COLCRYS</u>						
N 022352	001 7964648	Oct 06, 2028	U-1161			
	7981938	Oct 06, 2028	U-1166			
	8093296	Oct 06, 2028	U-1007			
	8093297	Oct 06, 2028	U-1161			
	8093298	Oct 06, 2028	U-1116			
	8097655	Oct 06, 2028	U-1020			
	8415395	Oct 06, 2028	U-1007			
	8415396	Oct 06, 2028	U-1007			
	8440721	Feb 17, 2029	U-1007			
	8440722	Feb 17, 2029	U-1020			
<u>COLCHICINE - MITIGARE</u>						
N 204820	001 8927607	Aug 22, 2033	U-1020			
	9399036	Aug 22, 2033	U-1020			
	9555029	Aug 22, 2033	U-1020			
	9675613	Aug 22, 2033	U-1020			
	9789108	Aug 22, 2033	U-1020			
<u>COLCHICINE - GLOPERBA</u>						
N 210942	001 10226423	Dec 20, 2037	DP			
	10383820	Nov 22, 2036	DP U-2814			
	10383821	Nov 22, 2036	DP			
	9907751	Nov 22, 2036	DP			
<u>COLCHICINE - LODOCO</u>						
N 215727	001 10130585	May 09, 2034	U-3642			
	10206891	Nov 01, 2033	U-3641			
	10265281	Nov 01, 2033	U-3639			
	10842762	Nov 01, 2033	U-3640			
	11026899	Nov 01, 2033	U-3639			
	11026900	Nov 01, 2033	U-3639			
	11026901	Nov 01, 2033	U-3638			
	11944594	Nov 01, 2033	U-3864			
	11944595	Nov 01, 2033	U-3867			
	11944595	Nov 01, 2033	U-3868			
	9744144	Nov 01, 2033	U-3643			
<u>COPANLISIB DIHYDROCHLORIDE - ALIOOPA</u>						
N 209936	001 10383876	Mar 29, 2032	DS DP			
	9636344	Mar 29, 2032	U-2124			
	RE46856	Oct 22, 2029	DS DP U-2124			
<u>COPPER - PARAGARD T 380A</u>						
N 018680	001				D-193	Jun 28, 2027
<u>COPPER CU-64 DOTATATE - DETECTNET</u>						
N 213227	001 10159759	Aug 23, 2032	U-2951		NCE	Sep 03, 2025
	10383961	Aug 23, 2032	U-2951		ODE-317	Sep 03, 2027
	11160888	Aug 23, 2032	U-2951			
	12102696	Sep 03, 2041	DP			
<u>CORTICOTROPIN - ACTHAR GEL (AUTOINJECTOR)</u>						
N 008372	003 11752199	Feb 25, 2041	U-3686			
	11752199	Feb 25, 2041	U-3687			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>CORTICOTROPIN - ACTHAR GEL (AUTOINJECTOR)</u>						
N 008372 003	11752199	Feb 25, 2041	U-3688			
<u>CORTICOTROPIN - ACTHAR GEL (AUTOINJECTOR)</u>						
N 008372 004	11752199	Feb 25, 2041	U-3686			
	11752199	Feb 25, 2041	U-3687			
	11752199	Feb 25, 2041	U-3688			
<u>CORTICOTROPIN - ACTHAR GEL</u>						
N 008372 008	11752199	Feb 25, 2041	U-3686			
	11752199	Feb 25, 2041	U-3687			
	11752199	Feb 25, 2041	U-3688			
<u>CORTICOTROPIN - PURIFIED CORTROPHIN GEL</u>						
N 008975 001	12102662	Oct 27, 2043	U-3904			
	12102662	Oct 27, 2043	U-3905			
	12102662	Oct 27, 2043	U-3906			
	12102662	Oct 27, 2043	U-3907			
	12102662	Oct 27, 2043	U-3908			
	12102662	Oct 27, 2043	U-3909			
	12102662	Oct 27, 2043	U-3910			
	12102662	Oct 27, 2043	U-3911			
	12102662	Oct 27, 2043	U-3912			
	12102662	Oct 27, 2043	U-3913			
	12102662	Oct 27, 2043	U-3914			
	12102662	Oct 27, 2043	U-3915			
	12102662	Oct 27, 2043	U-3916			
	12102662	Oct 27, 2043	U-3917			
	12102662	Oct 27, 2043	U-3918			
	12102662	Oct 27, 2043	U-3919			
	12102662	Oct 27, 2043	U-3920			
	12102662	Oct 27, 2043	U-3921			
	12102662	Oct 27, 2043	U-3922			
	12102662	Oct 27, 2043	U-3923			
	12102662	Oct 27, 2043	U-3924			
	12102662	Oct 27, 2043	U-3925			
	12102662	Oct 27, 2043	U-3926			
<u>CORTICOTROPIN - PURIFIED CORTROPHIN GEL</u>						
N 008975 002	11975047	Oct 27, 2043	U-3904			
	11975047	Oct 27, 2043	U-3905			
	11975047	Oct 27, 2043	U-3906			
	11975047	Oct 27, 2043	U-3907			
	11975047	Oct 27, 2043	U-3908			
	11975047	Oct 27, 2043	U-3909			
	11975047	Oct 27, 2043	U-3910			
	11975047	Oct 27, 2043	U-3911			
	11975047	Oct 27, 2043	U-3912			
	11975047	Oct 27, 2043	U-3913			
	11975047	Oct 27, 2043	U-3914			
	11975047	Oct 27, 2043	U-3915			
	11975047	Oct 27, 2043	U-3916			
	11975047	Oct 27, 2043	U-3917			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>CORTICOTROPIN - PURIFIED CORTROPHIN GEL</u>						
N 008975 002	11975047	Oct 27, 2043	U-3918			
	11975047	Oct 27, 2043	U-3919			
	11975047	Oct 27, 2043	U-3920			
	11975047	Oct 27, 2043	U-3921			
	11975047	Oct 27, 2043	U-3922			
	11975047	Oct 27, 2043	U-3923			
	11975047	Oct 27, 2043	U-3924			
	11975047	Oct 27, 2043	U-3925			
	11975047	Oct 27, 2043	U-3926			
	12102662	Oct 27, 2043	U-3904			
	12102662	Oct 27, 2043	U-3905			
	12102662	Oct 27, 2043	U-3906			
	12102662	Oct 27, 2043	U-3907			
	12102662	Oct 27, 2043	U-3908			
	12102662	Oct 27, 2043	U-3909			
	12102662	Oct 27, 2043	U-3910			
	12102662	Oct 27, 2043	U-3911			
	12102662	Oct 27, 2043	U-3912			
	12102662	Oct 27, 2043	U-3913			
	12102662	Oct 27, 2043	U-3914			
	12102662	Oct 27, 2043	U-3915			
	12102662	Oct 27, 2043	U-3916			
	12102662	Oct 27, 2043	U-3917			
	12102662	Oct 27, 2043	U-3918			
	12102662	Oct 27, 2043	U-3919			
	12102662	Oct 27, 2043	U-3920			
	12102662	Oct 27, 2043	U-3921			
	12102662	Oct 27, 2043	U-3922			
	12102662	Oct 27, 2043	U-3923			
	12102662	Oct 27, 2043	U-3924			
	12102662	Oct 27, 2043	U-3925			
	12102662	Oct 27, 2043	U-3926			
<u>CRINECERFONT - CRENESSITY</u>						
N 218808 001					ODE-503	Dec 13, 2031
<u>CRINECERFONT - CRENESSITY</u>						
N 218808 002					ODE-503	Dec 13, 2031
<u>CRINECERFONT - CRENESSITY</u>						
N 218808 003					ODE-503	Dec 13, 2031
<u>CRINECERFONT - CRENESSITY</u>						
N 218820 001					ODE-503	Dec 13, 2031
<u>CRISABOROLE - EUCRISA</u>						
N 207695 001	8039451	Jun 29, 2029	DS DP		D-191	Apr 03, 2026
	8039451*PED	Dec 29, 2029				
	8168614	Jan 20, 2030	U-1932			
	8168614*PED	Jul 20, 2030				
	8501712	Feb 16, 2027	U-1932			
	8501712*PED	Aug 16, 2027				
	9682092	Feb 16, 2027	U-1932			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>CRISABOROLE - EUCRISA</u>						
N 207695	001 9682092*PED	Aug 16, 2027				
<u>CRIZOTINIB - XALKORI</u>						
N 202570	001 7230098	Aug 26, 2025	DS		I-897	Jul 14, 2025
	7825137	May 12, 2027		U-3057	ODE-328	Jan 14, 2028
	7825137	May 12, 2027		U-3058	ODE-407	Jul 14, 2029
	7825137	May 12, 2027		U-3403		
	7858643	Oct 08, 2029	DS DP			
	8217057	Nov 06, 2029	DS DP			
	8785632	Mar 01, 2025	DS			
<u>CRIZOTINIB - XALKORI</u>						
N 202570	002 7230098	Aug 26, 2025	DS		I-897	Jul 14, 2025
	7825137	May 12, 2027		U-3057	ODE-328	Jan 14, 2028
	7825137	May 12, 2027		U-3058	ODE-407	Jul 14, 2029
	7825137	May 12, 2027		U-3403		
	7858643	Oct 08, 2029	DS DP			
	8217057	Nov 06, 2029	DS DP			
	8785632	Mar 01, 2025	DS			
<u>CRIZOTINIB - XALKORI</u>						
N 217581	001 7230098	Aug 26, 2025	DS		I-897	Jul 14, 2025
	7825137	May 12, 2027		U-3057		
	7825137	May 12, 2027		U-3058		
	7825137	May 12, 2027		U-3403		
	7858643	Oct 08, 2029	DS DP			
	8217057	Nov 06, 2029	DS			
	8785632	Mar 01, 2025	DS DP			
<u>CRIZOTINIB - XALKORI</u>						
N 217581	002 7230098	Aug 26, 2025	DS		I-897	Jul 14, 2025
	7825137	May 12, 2027		U-3057		
	7825137	May 12, 2027		U-3058		
	7825137	May 12, 2027		U-3403		
	7858643	Oct 08, 2029	DS DP			
	8217057	Nov 06, 2029	DS			
	8785632	Mar 01, 2025	DS DP			
<u>CRIZOTINIB - XALKORI</u>						
N 217581	003 7230098	Aug 26, 2025	DS		I-897	Jul 14, 2025
	7825137	May 12, 2027		U-3057		
	7825137	May 12, 2027		U-3058		
	7825137	May 12, 2027		U-3403		
	7858643	Oct 08, 2029	DS DP			
	8217057	Nov 06, 2029	DS			
	8785632	Mar 01, 2025	DS DP			
<u>CROFELEMER - MYTESI</u>						
N 202292	001 8962680	Oct 31, 2031		U-1319		
	9585868	Oct 31, 2031		U-1319		
<u>CUPRIC SULFATE; MANGANESE SULFATE; SELENIOUS ACID; ZINC SULFATE - TRALEMENT</u>						
N 209376	001 11786548	Jul 01, 2041		DP		
	11975022	Jul 01, 2041		U-3899		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>CUPRIC SULFATE; MANGANESE SULFATE; SELENIOS ACID; ZINC SULFATE - TRALEMENT</u>						
N 209376	001	11998565	Jul 01, 2041	DP		
		12150956	Jul 01, 2041	U-3899		
		12150957	Jul 01, 2041	U-3899		
<u>CUPRIC SULFATE; MANGANESE SULFATE; SELENIOS ACID; ZINC SULFATE - TRALEMENT</u>						
N 209376	002	11786548	Jul 01, 2041	DP		
		11975022	Jul 01, 2041	U-3899		
		11998565	Jul 01, 2041	DP		
		12150956	Jul 01, 2041	U-3899		
		12150957	Jul 01, 2041	U-3899		
<u>CUPRIC SULFATE; MANGANESE SULFATE; SELENIOS ACID; ZINC SULFATE - MULTRY5</u>						
N 209376	003	11786548	Jul 01, 2041	DP		
		11975022	Jul 01, 2041	U-3900		
		11998565	Jul 01, 2041	DP		
		12150956	Jul 01, 2041	U-3900		
		12150957	Jul 01, 2041	U-3900		
<u>CYCLOPHOSPHAMIDE - CYCLOPHOSPHAMIDE</u>						
N 210735	001	9662342	Jun 26, 2035	DP		
<u>CYCLOPHOSPHAMIDE - CYCLOPHOSPHAMIDE</u>						
N 210735	002	9662342	Jun 26, 2035	DP		
<u>CYCLOPHOSPHAMIDE - CYCLOPHOSPHAMIDE</u>						
N 210735	003	9662342	Jun 26, 2035	DP		
<u>CYCLOPHOSPHAMIDE - CYCLOPHOSPHAMIDE</u>						
N 210852	001	10849916	Jul 13, 2035	DP		
		11382923	Dec 01, 2035	DP		
<u>CYCLOPHOSPHAMIDE - CYCLOPHOSPHAMIDE</u>						
N 210852	002	10849916	Jul 13, 2035	DP		
		11382923	Dec 01, 2035	DP		
<u>CYCLOPHOSPHAMIDE - CYCLOPHOSPHAMIDE</u>						
N 210852	003	10849916	Jul 13, 2035	DP		
		11382923	Dec 01, 2035	DP		
<u>CYCLOPHOSPHAMIDE - CYCLOPHOSPHAMIDE</u>						
N 212501	001	10993952	Feb 15, 2036	DP		
<u>CYCLOPHOSPHAMIDE - CYCLOPHOSPHAMIDE</u>						
N 212501	002	10993952	Feb 15, 2036	DP		
<u>CYCLOPHOSPHAMIDE - CYCLOPHOSPHAMIDE</u>						
N 212501	003	10993952	Feb 15, 2036	DP		
<u>CYCLOSPORINE - RESTASIS MULTIDOSE</u>						
N 050790	002	8292129	Feb 25, 2031	DP		
		8561859	Apr 16, 2032	DP		
		9669974	May 11, 2034	DP		
		9676525	Feb 07, 2034	DP		
<u>CYCLOSPORINE - CEOUA</u>						
N 210913	001	10441630	Aug 23, 2033	DP		
		10918694	Feb 28, 2037	DP		
		11951153	Feb 28, 2037	U-1483		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>CYCLOSPORINE - CEQUA</u>						
N 210913	001 8980839	Aug 23, 2033	DP U-1483			
	9937225	Aug 23, 2033	DP U-1483			
<u>CYCLOSPORINE - VERKAZIA</u>						
N 214965	001 11612658	Jan 27, 2026	U-3560		ODE-358	Jun 23, 2028
	7973081	Jan 27, 2026	DP			
	8298568	Nov 03, 2027	DP			
	8524779	Jan 27, 2026	DP			
	9132071	Jun 02, 2029	DP			
	9220694	Jan 27, 2026	DP			
	9956289	Jan 27, 2026	DP			
<u>CYCLOSPORINE - VEVYE</u>						
N 217469	001 10813976	Sep 22, 2037	DP		NP	May 30, 2026
	11154513	Nov 20, 2038	DP U-1900			
	11413323	Oct 11, 2039	U-3627			
	12059449	Apr 01, 2042	DP U-1483			
	8614178	Dec 13, 2030	DP			
<u>CYSTEAMINE BITARTRATE - PROCYSBI</u>						
N 203389	001 10143665	Aug 16, 2036	U-1399		ODE-162	Dec 22, 2024
	10143665*PED	Feb 16, 2037			PED	Jun 22, 2025
	10328037	Aug 16, 2036	U-1399			
	10328037*PED	Feb 16, 2037				
	10548859	Aug 16, 2036	U-1399			
	10548859*PED	Feb 16, 2037				
	10905662	Aug 16, 2036	U-1399			
	10905662*PED	Feb 16, 2037				
	8026284	Sep 22, 2027	U-1399			
	8026284*PED	Mar 22, 2028				
	9173851	Jun 17, 2034	DP			
	9173851*PED	Dec 17, 2034				
	9192590	Jan 26, 2027	U-1399			
	9192590*PED	Jul 26, 2027				
	9198882	Jan 26, 2027	U-1399			
	9198882*PED	Jul 26, 2027				
	9233077	Jun 17, 2034	DP			
	9233077*PED	Dec 17, 2034				
	9925156	Jan 26, 2027	DS DP U-1399			
	9925156*PED	Jul 26, 2027				
	9925157	Jan 26, 2027	DS DP U-1399			
	9925157*PED	Jul 26, 2027				
	9925158	Jan 26, 2027	DS DP U-1399			
	9925158*PED	Jul 26, 2027				
<u>CYSTEAMINE BITARTRATE - PROCYSBI</u>						
N 203389	002 10143665	Aug 16, 2036	U-1399		ODE-162	Dec 22, 2024
	10143665*PED	Feb 16, 2037			PED	Jun 22, 2025
	10328037	Aug 16, 2036	U-1399			
	10328037*PED	Feb 16, 2037				
	10548859	Aug 16, 2036	U-1399			
	10548859*PED	Feb 16, 2037				

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>CYSTEAMINE BITARTRATE - PROCYSBI</u>						
N 203389	002	10905662	Aug 16, 2036	U-1399		
		10905662*PED	Feb 16, 2037			
		8026284	Sep 22, 2027	U-1399		
		8026284*PED	Mar 22, 2028			
		9173851	Jun 17, 2034	DP		
		9173851*PED	Dec 17, 2034			
		9192590	Jan 26, 2027	U-1399		
		9192590*PED	Jul 26, 2027			
		9198882	Jan 26, 2027	U-1399		
		9198882*PED	Jul 26, 2027			
		9233077	Jun 17, 2034	DP		
		9233077*PED	Dec 17, 2034			
		9925156	Jan 26, 2027	DS DP U-1399		
		9925156*PED	Jul 26, 2027			
		9925157	Jan 26, 2027	DS DP U-1399		
		9925157*PED	Jul 26, 2027			
		9925158	Jan 26, 2027	DS DP U-1399		
		9925158*PED	Jul 26, 2027			
<u>CYSTEAMINE BITARTRATE - PROCYSBI</u>						
N 213491	001	10143665	Aug 16, 2036	U-1399	ODE*	Dec 22, 2024
		10143665*PED	Feb 16, 2037		PED	Jun 22, 2025
		10328037	Aug 16, 2036	U-1399		
		10328037*PED	Feb 16, 2037			
		10548859	Aug 16, 2036	U-1399		
		10548859*PED	Feb 16, 2037			
		10905662	Aug 16, 2036	U-1399		
		10905662*PED	Feb 16, 2037			
		8026284	Sep 22, 2027	U-1399		
		8026284*PED	Mar 22, 2028			
		9173851	Jun 17, 2034	DP		
		9173851*PED	Dec 17, 2034			
		9192590	Jan 26, 2027	U-1399		
		9192590*PED	Jul 26, 2027			
		9198882	Jan 26, 2027	U-1399		
		9198882*PED	Jul 26, 2027			
		9233077	Jun 17, 2034	DP		
		9233077*PED	Dec 17, 2034			
		9925156	Jan 26, 2027	DP U-1399		
		9925156*PED	Jul 26, 2027			
		9925157	Jan 26, 2027	DP U-1399		
		9925157*PED	Jul 26, 2027			
		9925158	Jan 26, 2027	DP U-1399		
		9925158*PED	Jul 26, 2027			
<u>CYSTEAMINE BITARTRATE - PROCYSBI</u>						
N 213491	002	10143665	Aug 16, 2036	U-1399	ODE*	Dec 22, 2024
		10143665*PED	Feb 16, 2037		PED	Jun 22, 2025
		10328037	Aug 16, 2036	U-1399		
		10328037*PED	Feb 16, 2037			
		10548859	Aug 16, 2036	U-1399		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>CYSTEAMINE BITARTRATE - PROCYSBI</u>						
N 213491	002	10548859*PED	Feb 16, 2037			
		10905662	Aug 16, 2036	U-1399		
		10905662*PED	Feb 16, 2037			
		8026284	Sep 22, 2027	U-1399		
		8026284*PED	Mar 22, 2028			
		9173851	Jun 17, 2034	DP		
		9173851*PED	Dec 17, 2034			
		9192590	Jan 26, 2027	U-1399		
		9192590*PED	Jul 26, 2027			
		9198882	Jan 26, 2027	U-1399		
		9198882*PED	Jul 26, 2027			
		9233077	Jun 17, 2034	DP		
		9233077*PED	Dec 17, 2034			
		9925156	Jan 26, 2027	DP U-1399		
		9925156*PED	Jul 26, 2027			
		9925157	Jan 26, 2027	DP U-1399		
		9925157*PED	Jul 26, 2027			
		9925158	Jan 26, 2027	DP U-1399		
		9925158*PED	Jul 26, 2027			
<u>CYSTEINE HYDROCHLORIDE - ELCYS</u>						
N 210660	001	10478453	Jan 15, 2039	DP U-2752		
		10583155	Jan 15, 2039	DP U-2752		
		10653719	Jan 15, 2039	DP		
		10905713	Jan 15, 2039	DP		
		10905714	Jan 15, 2039	DP		
		10912795	Jan 15, 2039	DP		
		10918662	Jan 15, 2039	DP		
		10933089	Jan 15, 2039	DP		
		11510941	Jan 15, 2039	DP		
		11510942	Jan 15, 2039	DP U-2752		
		11642370	Jan 15, 2039	DP		
		11648262	Jan 15, 2039	DP		
		11679125	Jan 15, 2039	DP		
		11684636	Jan 15, 2039	DP		
		11826383	Jan 15, 2039	DP		
		11969439	Jan 15, 2039	U-2752		
<u>CYSTEINE HYDROCHLORIDE - NOURESS</u>						
N 212535	001	10478453	Jan 15, 2039	DP U-2752		
		10493051	Mar 15, 2039	DP		
		10543186	Mar 15, 2039	U-2722		
		10583155	Jan 15, 2039	DP U-2752		
		10653719	Jan 15, 2039	DP		
		10702490	Mar 15, 2039	DP		
		10905713	Jan 15, 2039	DP		
		10905714	Jan 15, 2039	DP		
		10912795	Jan 15, 2039	DP		
		10918662	Jan 15, 2039	DP		
		10933089	Jan 15, 2039	DP		
		11045438	Mar 15, 2039	DP		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>CYSTEINE HYDROCHLORIDE - NOURESS</u>						
N 212535	001	11510941	Mar 02, 2039	DP		
		11510942	Jan 15, 2039	DP	U-2752	
		11642370	Jan 15, 2039	DP		
		11648262	Jan 15, 2039	DP		
		11679125	Jan 15, 2039	DP		
		11684636	Jan 15, 2039	DP		
		11826383	Jan 15, 2039	DP		
<u>CYTARABINE; DAUNORUBICIN - VYXEOS</u>						
N 209401	001	10028912	Oct 15, 2032	DP	U-3149	ODE-350 Mar 30, 2028
		10028912	Oct 15, 2032	DP	U-3150	
		10166184	Oct 15, 2032	DP	U-3149	
		10835492	Oct 15, 2032		U-3150	
		7850990	Jan 23, 2027	DP	U-3147	
		8022279	Sep 14, 2027	DP	U-3147	
		8092828	Apr 01, 2029		U-3147	
		8431806	Apr 22, 2025	DP	U-3147	
		8518437	Jun 07, 2026	DP		
		9271931	Jan 23, 2027	DP		
<u>DABIGATRAN ETEXILATE MESYLATE - PRADAXA</u>						
N 022512	001	7932273	Sep 07, 2025	DS	DP	
		9034822	Jan 20, 2031		U-1759	
		9034822*PED	Jul 20, 2031			
<u>DABIGATRAN ETEXILATE MESYLATE - PRADAXA</u>						
N 022512	002	7932273	Sep 07, 2025	DS	DP	
		9034822	Jan 20, 2031		U-1759	
		9034822*PED	Jul 20, 2031			
<u>DABIGATRAN ETEXILATE MESYLATE - PRADAXA</u>						
N 022512	003	7932273	Sep 07, 2025	DS	DP	
		7932273*PED	Mar 07, 2026			
		9034822	Jan 20, 2031		U-1759	
		9034822*PED	Jul 20, 2031			
<u>DABIGATRAN ETEXILATE MESYLATE - PRADAXA</u>						
N 214358	001	7932273	Sep 07, 2025	DS	DP	
		7932273*PED	Mar 07, 2026			
<u>DABIGATRAN ETEXILATE MESYLATE - PRADAXA</u>						
N 214358	002	7932273	Sep 07, 2025	DS	DP	
		7932273*PED	Mar 07, 2026			
<u>DABIGATRAN ETEXILATE MESYLATE - PRADAXA</u>						
N 214358	003	7932273	Sep 07, 2025	DS	DP	
		7932273*PED	Mar 07, 2026			
<u>DABIGATRAN ETEXILATE MESYLATE - PRADAXA</u>						
N 214358	004	7932273	Sep 07, 2025	DS	DP	
		7932273*PED	Mar 07, 2026			
<u>DABIGATRAN ETEXILATE MESYLATE - PRADAXA</u>						
N 214358	005	7932273	Sep 07, 2025	DS	DP	
		7932273*PED	Mar 07, 2026			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>DABIGATRAN ETEXILATE MESYLATE - PRADAXA</u>						
N 214358	006 7932273	Sep 07, 2025	DS DP			
	7932273*PED	Mar 07, 2026				
<u>DABRAFENIB MESYLATE - TAFINLAR</u>						
N 202806	001 10869869	Aug 30, 2033		U-3185	I-894	Jun 22, 2025
	10869869*PED	Feb 28, 2034			I-908	Mar 16, 2026
	7994185	Jan 20, 2030	DS DP	U-1406	ODE-182	Apr 30, 2025
	7994185	Jan 20, 2030	DS DP	U-2031	ODE-183	May 04, 2025
	7994185	Jan 20, 2030	DS DP	U-2032	ODE-428	Mar 16, 2030
	7994185	Jan 20, 2030	DS DP	U-2296	PED	Oct 30, 2025
	7994185*PED	Jul 20, 2030			PED	Nov 04, 2025
	8415345	Jan 20, 2030	DS DP	U-1406	PED	Dec 22, 2025
	8415345	Jan 20, 2030	DS DP	U-2031	PED	Sep 16, 2026
	8415345	Jan 20, 2030	DS DP	U-2032	PED	Sep 16, 2030
	8415345	Jan 20, 2030	DS DP	U-2296		
	8415345*PED	Jul 20, 2030				
	8703781	Oct 15, 2030	DS DP	U-1713		
	8703781	Oct 15, 2030	DS DP	U-2032		
	8703781	Oct 15, 2030	DS DP	U-2296		
	8703781	Oct 15, 2030	DS DP	U-2298		
	8703781*PED	Apr 15, 2031				
	8835443	Jun 10, 2025		U-2026		
	8835443	Jun 10, 2025		U-2027		
	8835443	Jun 10, 2025		U-2296		
	8835443	Jun 10, 2025		U-2298		
	8835443*PED	Dec 10, 2025				
	8952018	Oct 15, 2030		U-2027		
	8952018*PED	Apr 15, 2031				
	9233956	May 04, 2029		U-1811		
	9233956	May 04, 2029		U-2031		
	9233956	May 04, 2029		U-2032		
	9233956	May 04, 2029		U-2296		
	9233956*PED	Nov 04, 2029				
<u>DABRAFENIB MESYLATE - TAFINLAR</u>						
N 202806	002 10869869	Aug 30, 2033		U-3185	I-894	Jun 22, 2025
	10869869*PED	Feb 28, 2034			I-908	Mar 16, 2026
	7994185	Jan 20, 2030	DS DP	U-1406	ODE-182	Apr 30, 2025
	7994185	Jan 20, 2030	DS DP	U-2031	ODE-183	May 04, 2025
	7994185	Jan 20, 2030	DS DP	U-2032	ODE-428	Mar 16, 2030
	7994185	Jan 20, 2030	DS DP	U-2296	PED	Oct 30, 2025
	7994185*PED	Jul 20, 2030			PED	Nov 04, 2025
	8415345	Jan 20, 2030	DS DP	U-1406	PED	Dec 22, 2025
	8415345	Jan 20, 2030	DS DP	U-2031	PED	Sep 16, 2026
	8415345	Jan 20, 2030	DS DP	U-2032	PED	Sep 16, 2030
	8415345	Jan 20, 2030	DS DP	U-2296		
	8415345*PED	Jul 20, 2030				
	8703781	Oct 15, 2030	DS DP	U-1713		
	8703781	Oct 15, 2030	DS DP	U-2032		
	8703781	Oct 15, 2030	DS DP	U-2296		
	8703781	Oct 15, 2030	DS DP	U-2298		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>DABRAFENIB MESYLATE - TAFINLAR</u>						
N 202806 002	8703781*PED	Apr 15, 2031				
	8835443	Jun 10, 2025	U-2026			
	8835443	Jun 10, 2025	U-2027			
	8835443	Jun 10, 2025	U-2296			
	8835443	Jun 10, 2025	U-2298			
	8835443*PED	Dec 10, 2025				
	8952018	Oct 15, 2030	U-2027			
	8952018*PED	Apr 15, 2031				
	9233956	May 04, 2029	U-1811			
	9233956	May 04, 2029	U-2031			
	9233956	May 04, 2029	U-2032			
	9233956	May 04, 2029	U-2296			
	9233956*PED	Nov 04, 2029				
<u>DABRAFENIB MESYLATE - TAFINLAR</u>						
N 217514 001	11504333	Jun 29, 2038	DP		NP	Mar 16, 2026
	11504333*PED	Dec 29, 2038			ODE-428	Mar 16, 2030
	7994185	Jan 20, 2030	DS DP		PED	Sep 16, 2026
	7994185*PED	Jul 20, 2030			PED	Sep 16, 2030
	8415345	Jan 20, 2030	DS DP			
	8415345*PED	Jul 20, 2030				
	8703781	Oct 15, 2030	DS DP U-3565			
	8703781*PED	Apr 15, 2031				
	8835443	Jun 10, 2025	U-3565			
	8835443*PED	Dec 10, 2025				
<u>DACLATASVIR DIHYDROCHLORIDE - DAKLINZA</u>						
N 206843 001	8329159	Jul 24, 2029	DS			
	8629171	Jun 13, 2031	DS DP U-1724			
	8642025	Aug 11, 2027	DS DP U-1724			
	8642025	Aug 11, 2027	DS DP U-1725			
	8900566	Aug 08, 2027	U-1724			
	8900566	Aug 08, 2027	U-1725			
	9421192	Aug 08, 2027	DS U-1724			
	9421192	Aug 08, 2027	DS U-1725			
<u>DACLATASVIR DIHYDROCHLORIDE - DAKLINZA</u>						
N 206843 002	8329159	Jul 24, 2029	DS			
	8629171	Jun 13, 2031	DS DP U-1724			
	8642025	Aug 11, 2027	DS DP U-1724			
	8642025	Aug 11, 2027	DS DP U-1725			
	8900566	Aug 08, 2027	U-1724			
	8900566	Aug 08, 2027	U-1725			
	9421192	Aug 08, 2027	DS U-1724			
	9421192	Aug 08, 2027	DS U-1725			
<u>DACLATASVIR DIHYDROCHLORIDE - DAKLINZA</u>						
N 206843 003	9421192	Aug 08, 2027	DS U-1724			
	9421192	Aug 08, 2027	DS U-1725			
<u>DACOMITINIB - VIZIMPRO</u>						
N 211288 001	10596162	Feb 02, 2026	U-3338		ODE-206	Sep 27, 2025
	10603314	Feb 02, 2026	U-3337		ODE-213	Sep 27, 2025

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>DACOMITINIB - VIZIMPRO</u>						
N 211288	001 7772243	Aug 26, 2028	DS DP			
	8623883	May 05, 2025		U-1403		
<u>DACOMITINIB - VIZIMPRO</u>						
N 211288	002 10596162	Feb 02, 2026		U-3338	ODE-206	Sep 27, 2025
	10603314	Feb 02, 2026		U-3337	ODE-213	Sep 27, 2025
	7772243	Aug 26, 2028	DS DP			
	8623883	May 05, 2025		U-1403		
<u>DACOMITINIB - VIZIMPRO</u>						
N 211288	003 10596162	Feb 02, 2026		U-3338	ODE-206	Sep 27, 2025
	10603314	Feb 02, 2026		U-3337	ODE-213	Sep 27, 2025
	7772243	Aug 26, 2028	DS DP			
	8623883	May 05, 2025		U-1403		
<u>DALBAVANCIN HYDROCHLORIDE - DALVANCE</u>						
N 021883	001 6900175	May 23, 2028		U-3499		
<u>DANICOPAN - VOYDEYA</u>						
N 218037	001 12076319	Aug 02, 2038		U-3933	NCE	Mar 29, 2029
	9796741	Feb 25, 2035	DS	U-3933	ODE-476	Mar 29, 2031
<u>DANICOPAN - VOYDEYA</u>						
N 218037	002 12076319	Aug 02, 2038		U-3933	NCE	Mar 29, 2029
	9796741	Feb 25, 2035	DS	U-3933	ODE-476	Mar 29, 2031
<u>DANTROLENE SODIUM - RYANODEX</u>						
N 205579	001 7758890	Jun 30, 2025	DP			
<u>DAPAGLIFLOZIN - FARXIGA</u>						
N 202293	001 6515117	Oct 04, 2025	DS DP	U-2139	M-298	May 08, 2026
	6515117	Oct 04, 2025	DS DP	U-493	NPP	Jun 12, 2027
	6515117*PED	Apr 04, 2026			PED	Nov 08, 2026
	7456254	Jun 30, 2025		U-2139	PED	Dec 12, 2027
	7456254*PED	Dec 30, 2025				
	7851502	Aug 19, 2028	DP			
	7851502*PED	Feb 19, 2029				
	7919598	Dec 16, 2029	DS			
	7919598*PED	Jun 16, 2030				
	8221786	Mar 21, 2028	DP			
	8221786*PED	Sep 21, 2028				
	8329648	Aug 18, 2026		U-2139		
	8329648	Aug 18, 2026		U-2212		
	8329648	Aug 18, 2026		U-2213		
	8329648*PED	Feb 18, 2027				
	8361972	Mar 21, 2028		U-2139		
	8361972	Mar 21, 2028		U-493		
	8361972*PED	Sep 21, 2028				
	8431685	Apr 13, 2025		U-2139		
	8431685*PED	Oct 13, 2025				
	8461105	Apr 13, 2025		U-2139		
	8461105*PED	Oct 13, 2025				
	8501698	Jun 20, 2027	DP	U-493		
	8501698*PED	Dec 20, 2027				

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>DAPAGLIFLOZIN - FARXIGA</u>						
N 202293 001	8685934	May 26, 2030	U-1522			
	8685934*PED	Nov 26, 2030				
	8716251	Mar 21, 2028	DP			
	8716251*PED	Sep 21, 2028				
	8721615	Jan 18, 2030	DP	Y		
	8721615*PED	Jul 18, 2030				
	8906851	Aug 18, 2026	U-2139			
	8906851*PED	Feb 18, 2027				
<u>DAPAGLIFLOZIN - FARXIGA</u>						
N 202293 002	10973836	Mar 09, 2040	U-3127		M-298	May 08, 2026
	10973836*PED	Sep 09, 2040			NPP	Jun 12, 2027
	11826376	Jul 18, 2039	U-3766		PED	Nov 08, 2026
	11826376*PED	Jan 18, 2040			PED	Dec 12, 2027
	11903955	Mar 09, 2040	U-3825			
	11903955*PED	Sep 09, 2040				
	6515117	Oct 04, 2025	DS DP U-2139			
	6515117	Oct 04, 2025	DS DP U-493			
	6515117*PED	Apr 04, 2026				
	7456254	Jun 30, 2025	U-2139			
	7456254*PED	Dec 30, 2025				
	7851502	Aug 19, 2028	DP			
	7851502*PED	Feb 19, 2029				
	7919598	Dec 16, 2029	DS			
	7919598*PED	Jun 16, 2030				
	8221786	Mar 21, 2028	DP			
	8221786*PED	Sep 21, 2028				
	8329648	Aug 18, 2026	U-2139			
	8329648	Aug 18, 2026	U-2212			
	8329648	Aug 18, 2026	U-2213			
	8329648*PED	Feb 18, 2027				
	8361972	Mar 21, 2028	U-2139			
	8361972	Mar 21, 2028	U-493			
	8361972*PED	Sep 21, 2028				
	8431685	Apr 13, 2025	U-2139			
	8431685*PED	Oct 13, 2025				
	8461105	Apr 13, 2025	U-2139			
	8461105*PED	Oct 13, 2025				
	8501698	Jun 20, 2027	DP U-493			
	8501698*PED	Dec 20, 2027				
	8685934	May 26, 2030	U-1522			
	8685934*PED	Nov 26, 2030				
	8716251	Mar 21, 2028	DP			
	8716251*PED	Sep 21, 2028				
	8721615	Jan 18, 2030	DP	Y		
	8721615*PED	Jul 18, 2030				
	8906851	Aug 18, 2026	U-2139			
	8906851*PED	Feb 18, 2027				

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>DAPAGLIFLOZIN; METFORMIN HYDROCHLORIDE - XIGDUO XR</u>						
N 205649 001	6515117	Oct 04, 2025	DS DP U-493		NPP	Jun 12, 2027
	6515117*PED	Apr 04, 2026			PED	Dec 12, 2027
	7919598	Dec 16, 2029	DS			
	7919598*PED	Jun 16, 2030				
	8501698	Jun 20, 2027	DP U-493			
	8501698*PED	Dec 20, 2027				
	8685934	May 26, 2030	U-1522			
	8685934*PED	Nov 26, 2030				
	9616028	Nov 12, 2030	DP			
	9616028*PED	May 12, 2031				
<u>DAPAGLIFLOZIN; METFORMIN HYDROCHLORIDE - XIGDUO XR</u>						
N 205649 002	6515117	Oct 04, 2025	DS DP U-493		NPP	Jun 12, 2027
	6515117*PED	Apr 04, 2026			PED	Dec 12, 2027
	7919598	Dec 16, 2029	DS			
	7919598*PED	Jun 16, 2030				
	8501698	Jun 20, 2027	DP U-493			
	8501698*PED	Dec 20, 2027				
	8685934	May 26, 2030	U-1522			
	8685934*PED	Nov 26, 2030				
	9616028	Nov 12, 2030	DP			
	9616028*PED	May 12, 2031				
<u>DAPAGLIFLOZIN; METFORMIN HYDROCHLORIDE - XIGDUO XR</u>						
N 205649 003	6515117	Oct 04, 2025	DS DP U-493		NPP	Jun 12, 2027
	6515117*PED	Apr 04, 2026			PED	Dec 12, 2027
	7919598	Dec 16, 2029	DS			
	7919598*PED	Jun 16, 2030				
	8501698	Jun 20, 2027	DP U-493			
	8501698*PED	Dec 20, 2027				
	8685934	May 26, 2030	U-1522			
	8685934*PED	Nov 26, 2030				
	9616028	Nov 12, 2030	DP			
	9616028*PED	May 12, 2031				
<u>DAPAGLIFLOZIN; METFORMIN HYDROCHLORIDE - XIGDUO XR</u>						
N 205649 004	6515117	Oct 04, 2025	DS DP U-493		NPP	Jun 12, 2027
	6515117*PED	Apr 04, 2026			PED	Dec 12, 2027
	7919598	Dec 16, 2029	DS			
	7919598*PED	Jun 16, 2030				
	8501698	Jun 20, 2027	DP U-493			
	8501698*PED	Dec 20, 2027				
	8685934	May 26, 2030	U-1522			
	8685934*PED	Nov 26, 2030				
	9616028	Nov 12, 2030	DP			
	9616028*PED	May 12, 2031				
<u>DAPAGLIFLOZIN; METFORMIN HYDROCHLORIDE - XIGDUO XR</u>						
N 205649 005	6515117	Oct 04, 2025	DS DP U-493		NPP	Jun 12, 2027
	6515117*PED	Apr 04, 2026			PED	Dec 12, 2027
	7919598	Dec 16, 2029	DS			
	7919598*PED	Jun 16, 2030				

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>DAPAGLIFLOZIN; METFORMIN HYDROCHLORIDE - XIGDUO XR</u>						
N 205649 005	8501698	Jun 20, 2027	DP U-493			
	8501698*PED	Dec 20, 2027				
	8685934	May 26, 2030	U-1522			
	8685934*PED	Nov 26, 2030				
	9616028	Nov 12, 2030	DP			
	9616028*PED	May 12, 2031				
<u>DAPAGLIFLOZIN; METFORMIN HYDROCHLORIDE; SAXAGLIPTIN HYDROCHLORIDE - QTERNMET XR</u>						
N 210874 001	6515117	Oct 04, 2025	DS DP U-493			
	6515117*PED	Apr 04, 2026				
	7919598	Dec 16, 2029	DS			
	7919598*PED	Jun 16, 2030				
	8501698	Jun 20, 2027	DP U-493			
	8501698*PED	Dec 20, 2027				
	8628799	Jul 13, 2025	DP			
	8716251	Mar 21, 2028	DP			
	8716251*PED	Sep 21, 2028				
	9616028	Nov 12, 2030	DP			
	9616028*PED	May 12, 2031				
<u>DAPAGLIFLOZIN; METFORMIN HYDROCHLORIDE; SAXAGLIPTIN HYDROCHLORIDE - QTERNMET XR</u>						
N 210874 002	6515117	Oct 04, 2025	DS DP U-493			
	6515117*PED	Apr 04, 2026				
	7919598	Dec 16, 2029	DS			
	7919598*PED	Jun 16, 2030				
	8501698	Jun 20, 2027	DP U-493			
	8501698*PED	Dec 20, 2027				
	8628799	Jul 13, 2025	DP			
	8716251	Mar 21, 2028	DP			
	8716251*PED	Sep 21, 2028				
	9616028	Nov 12, 2030	DP			
	9616028*PED	May 12, 2031				
<u>DAPAGLIFLOZIN; METFORMIN HYDROCHLORIDE; SAXAGLIPTIN HYDROCHLORIDE - QTERNMET XR</u>						
N 210874 003	6515117	Oct 04, 2025	DS DP U-493			
	6515117*PED	Apr 04, 2026				
	7919598	Dec 16, 2029	DS			
	7919598*PED	Jun 16, 2030				
	8501698	Jun 20, 2027	DP U-493			
	8501698*PED	Dec 20, 2027				
	8628799	Jul 13, 2025	DP			
	8716251	Mar 21, 2028	DP			
	8716251*PED	Sep 21, 2028				
	9616028	Nov 12, 2030	DP			
	9616028*PED	May 12, 2031				
<u>DAPAGLIFLOZIN; METFORMIN HYDROCHLORIDE; SAXAGLIPTIN HYDROCHLORIDE - QTERNMET XR</u>						
N 210874 004	6515117	Oct 04, 2025	DS DP U-493			
	6515117*PED	Apr 04, 2026				
	7919598	Dec 16, 2029	DS			
	7919598*PED	Jun 16, 2030				
	8501698	Jun 20, 2027	DP U-493			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>DAPAGLIFLOZIN; METFORMIN HYDROCHLORIDE; SAXAGLIPTIN HYDROCHLORIDE - QTERNMET XR</u>						
N 210874	004	8501698*PED	Dec 20, 2027			
		8628799	Jul 13, 2025	DP		
		8716251	Mar 21, 2028	DP		
		8716251*PED	Sep 21, 2028			
		9616028	Nov 12, 2030	DP		
		9616028*PED	May 12, 2031			
<u>DAPAGLIFLOZIN; SAXAGLIPTIN HYDROCHLORIDE - QTERN</u>						
N 209091	001	6515117	Oct 04, 2025	DS DP U-493		
		6515117*PED	Apr 04, 2026			
		7919598	Dec 16, 2029	DS		
		7919598*PED	Jun 16, 2030			
		8221786	Mar 21, 2028	DP		
		8221786*PED	Sep 21, 2028			
		8361972	Mar 21, 2028		U-1976	
		8361972	Mar 21, 2028		U-1977	
		8361972	Mar 21, 2028		U-493	
		8361972*PED	Sep 21, 2028			
		8501698	Jun 20, 2027	DP U-1976		
		8501698	Jun 20, 2027	DP U-1977		
		8501698	Jun 20, 2027	DP U-493		
		8501698*PED	Dec 20, 2027			
		8628799	Jul 13, 2025	DP		
		8716251	Mar 21, 2028	DP		
		8716251*PED	Sep 21, 2028			
<u>DAPAGLIFLOZIN; SAXAGLIPTIN HYDROCHLORIDE - QTERN</u>						
N 209091	002	6515117	Oct 04, 2025	DS DP U-493		
		6515117*PED	Apr 04, 2026			
		7919598	Dec 16, 2029	DS		
		7919598*PED	Jun 16, 2030			
		8221786	Mar 21, 2028	DP		
		8221786*PED	Sep 21, 2028			
		8361972	Mar 21, 2028		U-493	
		8361972*PED	Sep 21, 2028			
		8501698	Jun 20, 2027	DP U-493		
		8501698*PED	Dec 20, 2027			
		8628799	Jul 13, 2025	DP		
		8716251	Mar 21, 2028	DP		
		8716251*PED	Sep 21, 2028			
<u>DAPRODUSTAT - JESDUVROO</u>						
N 216951	001	11117871	Mar 13, 2038	DS DP U-3535	NCE	Feb 01, 2028
		11643397	Jun 22, 2027	DS U-3535		
		11649217	Mar 13, 2038	DP U-3535		
		8324208	Dec 11, 2028	DS DP		
		8557834	Jun 22, 2027		U-1238	
		8815884	Jun 22, 2027	DP U-1238		
<u>DAPRODUSTAT - JESDUVROO</u>						
N 216951	002	11117871	Mar 13, 2038	DS DP U-3535	NCE	Feb 01, 2028
		11643397	Jun 22, 2027	DS U-3535		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>DAPRODUSTAT - JESDUVROO</u>						
N 216951 002	11649217	Mar 13, 2038	DP U-3535			
	8324208	Dec 11, 2028	DS DP			
	8557834	Jun 22, 2027	U-1238			
	8815884	Jun 22, 2027	DP U-1238			
<u>DAPRODUSTAT - JESDUVROO</u>						
N 216951 003	11117871	Mar 13, 2038	DS DP U-3535		NCE	Feb 01, 2028
	11643397	Jun 22, 2027	DS U-3535			
	11649217	Mar 13, 2038	DP U-3535			
	8324208	Dec 11, 2028	DS DP			
	8557834	Jun 22, 2027	U-1238			
	8815884	Jun 22, 2027	DP U-1238			
<u>DAPRODUSTAT - JESDUVROO</u>						
N 216951 004	11117871	Mar 13, 2038	DS DP U-3535		NCE	Feb 01, 2028
	11643397	Jun 22, 2027	DS U-3535			
	11649217	Mar 13, 2038	DP U-3535			
	8324208	Dec 11, 2028	DS DP			
	8557834	Jun 22, 2027	U-1238			
	8815884	Jun 22, 2027	DP U-1238			
<u>DAPRODUSTAT - JESDUVROO</u>						
N 216951 005	11117871	Mar 13, 2038	DS DP U-3535		NCE	Feb 01, 2028
	11643397	Jun 22, 2027	DS U-3535			
	11649217	Mar 13, 2038	DP U-3535			
	8324208	Dec 11, 2028	DS DP			
	8557834	Jun 22, 2027	U-1238			
	8815884	Jun 22, 2027	DP U-1238			
<u>DAPSONE - ACZONE</u>						
N 207154 001	11273132	Nov 18, 2033	DP			
<u>DAPTOMYCIN - CUBICIN</u>						
N 021572 002	8003673	Sep 04, 2028	U-1180			
<u>DAPTOMYCIN - CUBICIN RF</u>						
N 021572 003	9138456	Nov 23, 2030	DP			
<u>DAPTOMYCIN - DAPTOMYCIN</u>						
N 210282 001	10357535	Sep 11, 2033	DP U-3176			
	9655946	Sep 11, 2033	DP U-3175			
<u>DAPTOMYCIN - DAPTOMYCIN</u>						
N 210282 002	10357535	Sep 11, 2033	DP U-3176			
	9655946	Sep 11, 2033	DP U-3175			
<u>DAPTOMYCIN - DAPZURA RT</u>						
N 213645 001	11173189	Mar 11, 2041	DP U-3294			
<u>DAPTOMYCIN - DAPTOMYCIN</u>						
N 217415 001	11759497	Aug 28, 2038	DP			
	12053502	Aug 28, 2038	U-3981			
<u>DAPTOMYCIN - DAPTOMYCIN</u>						
N 217415 002	11759497	Aug 28, 2038	DP			
	12053502	Aug 28, 2038	U-3981			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>DARIDOREXANT HYDROCHLORIDE - QUVIVIQ</u>						
N 214985 001	10023560	Dec 02, 2034	U-620		M-310	Sep 30, 2027
	9732075	Jun 12, 2033	DS DP U-620		NCE	Apr 07, 2027
	9790208	Dec 02, 2034	DS DP			
<u>DARIDOREXANT HYDROCHLORIDE - QUVIVIQ</u>						
N 214985 002	10023560	Dec 02, 2034	U-620		M-310	Sep 30, 2027
	9732075	Jun 12, 2033	DS DP U-620		NCE	Apr 07, 2027
	9790208	Dec 02, 2034	DS DP			
<u>DAROLUTAMIDE - NUBEQA</u>						
N 212099 001	10010530	Jan 28, 2036	DS		I-900	Aug 05, 2025
	10383853	Jan 28, 2036	DS			
	10711013	Oct 27, 2030	DS DP			
	10835515	Jan 28, 2036	DP U-2605			
	11046713	Oct 27, 2030	DS			
	11168058	Feb 27, 2038	DS DP			
	8975254	Mar 25, 2033	DS DP U-2605			
	9657003	Oct 27, 2030	DS DP U-2605			
<u>DARUNAVIR - PREZISTA</u>						
N 021976 001	7700645	Dec 26, 2026	DS DP			
<u>DARUNAVIR - PREZISTA</u>						
N 021976 002	7700645	Dec 26, 2026	DS DP			
<u>DARUNAVIR - PREZISTA</u>						
N 021976 003	7700645	Dec 26, 2026	DS DP			
<u>DARUNAVIR - PREZISTA</u>						
N 021976 004	7700645	Dec 26, 2026	DS DP			
<u>DARUNAVIR - PREZISTA</u>						
N 021976 005	7700645	Dec 26, 2026	DS DP			
<u>DARUNAVIR - PREZISTA</u>						
N 021976 006	7700645	Dec 26, 2026	DS DP			
<u>DARUNAVIR - PREZISTA</u>						
N 202895 001	7700645	Dec 26, 2026	DS DP			
<u>DASABUVIR SODIUM; OMBITASVIR, PARITAPREVIR, RITONAVIR - VIEKIRA PAK (COPACKAGED)</u>						
N 206619 001	10201542	Oct 18, 2033	DP U-1753			
	8188104	May 17, 2029	DS DP U-1636			
	8420596	Apr 10, 2031	DS DP			
	8466159	Sep 04, 2032	U-1637			
	8492386	Sep 04, 2032	U-1840			
	8501238	Dec 19, 2028	DS DP U-1636			
	8642538	Sep 10, 2029	DS DP U-1638			
	8680106	Sep 04, 2032	U-1637			
	8685984	Sep 04, 2032	U-1840			
	8686026	Jun 09, 2031	DP			
	8691938	Apr 13, 2032	DS DP			
	9006387	Jun 10, 2030	U-1687			
	9044480	Apr 10, 2031	U-1638			
	9139536	Nov 09, 2028	U-1753			
	9629841	Oct 18, 2033	DP U-1753			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>DASABUVIR SODIUM; OMBITASVIR, PARITAPREVIR, RITONAVIR - VIEKIRA PAK (COPACKAGED)</u>						
N 206619 001	10201542	Oct 18, 2033	DP U-1753			
	8188104	May 17, 2029	DS DP U-1636			
	8420596	Apr 10, 2031	DS DP			
	8466159	Sep 04, 2032	U-1637			
	8492386	Sep 04, 2032	U-1840			
	8501238	Dec 19, 2028	DS DP U-1636			
	8642538	Sep 10, 2029	DS DP U-1638			
	8680106	Sep 04, 2032	U-1637			
	8685984	Sep 04, 2032	U-1840			
	8686026	Jun 09, 2031	DP			
	8691938	Apr 13, 2032	DS DP			
	9006387	Jun 10, 2030	U-1687			
	9044480	Apr 10, 2031	U-1638			
	9139536	Nov 09, 2028	U-1753			
	9629841	Oct 18, 2033	DP U-1753			
<u>DASABUVIR SODIUM; OMBITASVIR; PARITAPREVIR; RITONAVIR - VIEKIRA XR</u>						
N 208624 001	10105365	Jan 02, 2035	DP U-1889			
	10201541	May 17, 2032	DP			
	10201584	May 17, 2032	U-1889			
	8188104	May 17, 2029	DS DP U-1636			
	8420596	Apr 10, 2031	DS DP			
	8466159	Sep 04, 2032	U-1637			
	8492386	Sep 04, 2032	U-1840			
	8501238	Sep 17, 2028	DS DP U-1636			
	8642538	Sep 10, 2029	DS DP U-1638			
	8680106	Sep 04, 2032	U-1637			
	8685984	Sep 04, 2032	U-1840			
	8686026	Jun 09, 2031	DP			
	8691938	Apr 13, 2032	DS DP			
	9006387	Jun 10, 2030	U-1687			
	9044480	Apr 10, 2031	U-1638			
	9139536	Nov 09, 2028	U-1753			
	9333204	Jan 02, 2035	DP U-1889			
	9744170	Jan 02, 2035	DP U-1889			
<u>DASATINIB - DASATINIB</u>						
A 202103 001					PC	Mar 02, 2025
<u>DASATINIB - DASATINIB</u>						
A 202103 002					PC	Mar 02, 2025
<u>DASATINIB - DASATINIB</u>						
A 202103 003					PC	Mar 02, 2025
<u>DASATINIB - DASATINIB</u>						
A 202103 004					PC	Mar 02, 2025
<u>DASATINIB - DASATINIB</u>						
A 203180 001					PC	Mar 02, 2025
<u>DASATINIB - DASATINIB</u>						
A 203180 002					PC	Mar 02, 2025

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>DASATINIB - SPRYCEL</u>						
N 021986 001	7491725	Mar 28, 2026	DS DP		ODE-164	Nov 09, 2024
	7491725*PED	Sep 28, 2026			ODE-225	Dec 21, 2025
	8680103	Feb 04, 2025	DP		PED	May 09, 2025
	8680103*PED	Aug 04, 2025			PED	Jun 21, 2026
<u>DASATINIB - SPRYCEL</u>						
N 021986 002	7491725	Mar 28, 2026	DS DP		ODE-164	Nov 09, 2024
	7491725*PED	Sep 28, 2026			ODE-225	Dec 21, 2025
	8680103	Feb 04, 2025	DP		PED	May 09, 2025
	8680103*PED	Aug 04, 2025			PED	Jun 21, 2026
<u>DASATINIB - SPRYCEL</u>						
N 021986 003	7491725	Mar 28, 2026	DS DP		ODE-164	Nov 09, 2024
	7491725*PED	Sep 28, 2026			ODE-225	Dec 21, 2025
	8680103	Feb 04, 2025	DP		PED	May 09, 2025
	8680103*PED	Aug 04, 2025			PED	Jun 21, 2026
<u>DASATINIB - SPRYCEL</u>						
N 021986 004	7491725	Mar 28, 2026	DS DP		ODE-164	Nov 09, 2024
	7491725*PED	Sep 28, 2026			ODE-225	Dec 21, 2025
	8680103	Feb 04, 2025	DP		PED	May 09, 2025
	8680103*PED	Aug 04, 2025			PED	Jun 21, 2026
<u>DASATINIB - SPRYCEL</u>						
N 021986 005	7491725	Mar 28, 2026	DS DP		ODE-164	Nov 09, 2024
	7491725*PED	Sep 28, 2026			ODE-225	Dec 21, 2025
	8680103	Feb 04, 2025	DP		PED	May 09, 2025
	8680103*PED	Aug 04, 2025			PED	Jun 21, 2026
<u>DASATINIB - SPRYCEL</u>						
N 021986 006	7491725	Mar 28, 2026	DS DP		ODE-164	Nov 09, 2024
	7491725*PED	Sep 28, 2026			ODE-225	Dec 21, 2025
	8680103	Feb 04, 2025	DP		PED	May 09, 2025
	8680103*PED	Aug 04, 2025			PED	Jun 21, 2026
<u>DASATINIB - PHYRAGO</u>						
N 216099 001	11202778	Jan 22, 2041		U-3770	M-307	Dec 05, 2026
	11202778	Jan 22, 2041		U-3771		
	11202778	Jan 22, 2041		U-3772		
	11298356	Jan 22, 2041	DP			
	11324745	Jan 22, 2041		U-3767		
	11324745	Jan 22, 2041		U-3768		
	11324745	Jan 22, 2041		U-3769		
<u>DASATINIB - PHYRAGO</u>						
N 216099 002	11202778	Jan 22, 2041		U-3770	M-307	Dec 05, 2026
	11202778	Jan 22, 2041		U-3771		
	11202778	Jan 22, 2041		U-3772		
	11298356	Jan 22, 2041	DP			
	11324745	Jan 22, 2041		U-3767		
	11324745	Jan 22, 2041		U-3768		
	11324745	Jan 22, 2041		U-3769		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>DASATINIB - PHYRAGO</u>						
N 216099 003	11202778	Jan 22, 2041	U-3770		M-307	Dec 05, 2026
	11202778	Jan 22, 2041	U-3771			
	11202778	Jan 22, 2041	U-3772			
	11298356	Jan 22, 2041	DP			
	11324745	Jan 22, 2041	U-3767			
	11324745	Jan 22, 2041	U-3768			
	11324745	Jan 22, 2041	U-3769			
<u>DASATINIB - PHYRAGO</u>						
N 216099 004	11202778	Jan 22, 2041	U-3770		M-307	Dec 05, 2026
	11202778	Jan 22, 2041	U-3771			
	11202778	Jan 22, 2041	U-3772			
	11298356	Jan 22, 2041	DP			
	11324745	Jan 22, 2041	U-3767			
	11324745	Jan 22, 2041	U-3768			
	11324745	Jan 22, 2041	U-3769			
<u>DASATINIB - PHYRAGO</u>						
N 216099 005	11202778	Jan 22, 2041	U-3770		M-307	Dec 05, 2026
	11202778	Jan 22, 2041	U-3771			
	11202778	Jan 22, 2041	U-3772			
	11298356	Jan 22, 2041	DP			
	11324745	Jan 22, 2041	U-3767			
	11324745	Jan 22, 2041	U-3768			
	11324745	Jan 22, 2041	U-3769			
<u>DASATINIB - PHYRAGO</u>						
N 216099 006	11202778	Jan 22, 2041	U-3770		M-307	Dec 05, 2026
	11202778	Jan 22, 2041	U-3771			
	11202778	Jan 22, 2041	U-3772			
	11298356	Jan 22, 2041	DP			
	11324745	Jan 22, 2041	U-3767			
	11324745	Jan 22, 2041	U-3768			
	11324745	Jan 22, 2041	U-3769			
<u>DASIGLUCAGON HYDROCHLORIDE - ZEGALOGUE</u>						
N 214231 001	10442847	Feb 03, 2035	DS DP		NCE	Mar 22, 2026
	11795204	Jan 06, 2034	U-3752			
<u>DASIGLUCAGON HYDROCHLORIDE - ZEGALOGUE (AUTOINJECTOR)</u>						
N 214231 002	10442847	Feb 03, 2035	DS DP		NCE	Mar 22, 2026
	11795204	Jan 06, 2034	U-3752			
<u>DEFERASIROX - JADENU</u>						
N 206910 001	9283209	Nov 21, 2034	DS DP			
<u>DEFERASIROX - JADENU</u>						
N 206910 002	9283209	Nov 21, 2034	DS DP			
<u>DEFERASIROX - JADENU</u>						
N 206910 003	9283209	Nov 21, 2034	DS DP			
<u>DEFERIPRONE - FERRIPROX</u>						
N 021825 001					ODE-417	Apr 30, 2028
					ODE-420	Apr 30, 2028

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>DEFERIPRONE - FERRIPROX</u>						
N 021825	001				ODE-421	Apr 30, 2028
<u>DEFERIPRONE - FERRIPROX</u>						
N 021825	002				ODE-417	Apr 30, 2028
					ODE-420	Apr 30, 2028
					ODE-421	Apr 30, 2028
<u>DEFERIPRONE - FERRIPROX</u>						
N 208030	001	8703156	Oct 26, 2029	DP U-3083	ODE-417	Apr 30, 2028
					ODE-418	Apr 30, 2028
					ODE-419	Apr 30, 2028
<u>DEFERIPRONE - FERRIPROX</u>						
N 208030	002	8703156	Oct 26, 2029	DP U-3083	ODE-417	Apr 30, 2028
					ODE-418	Apr 30, 2028
					ODE-419	Apr 30, 2028
<u>DEFERIPRONE - FERRIPROX</u>						
N 212269	001	10780055	Oct 25, 2038	DP U-3083	ODE-417	Apr 30, 2028
		10940115	Oct 25, 2038	DP U-3083	ODE-420	Apr 30, 2028
		10940116	Oct 25, 2038	DP	ODE-421	Apr 30, 2028
		11357731	Oct 25, 2038	U-3083		
		11458103	Oct 25, 2038	DP		
		11723874	Oct 25, 2038	U-3083		
<u>DEFIBROTIDE SODIUM - DEFITELIO</u>						
N 208114	001	11085043	Jun 22, 2032	DP		
		11236328	Jun 22, 2032	U-3301		
		11746348	Jun 22, 2032	DP		
<u>DEFLAZACORT - EMFLAZA</u>						
N 208684	001				ODE-252	Jun 07, 2026
<u>DEFLAZACORT - EMFLAZA</u>						
N 208684	002				ODE-252	Jun 07, 2026
<u>DEFLAZACORT - EMFLAZA</u>						
N 208684	003				ODE-252	Jun 07, 2026
<u>DEFLAZACORT - EMFLAZA</u>						
N 208684	004				ODE-252	Jun 07, 2026
<u>DEFLAZACORT - EMFLAZA</u>						
N 208685	001				ODE-252	Jun 07, 2026
<u>DEGARELIX ACETATE - FIRMAGON</u>						
N 022201	001	10695398	Apr 27, 2032	U-1895		
		10729739	Feb 10, 2029	U-1978		
		10973870	Feb 10, 2029	U-1978		
		11766468	Apr 27, 2032	U-1978		
		11826397	Apr 27, 2032	U-1978		
		9415085	Apr 27, 2032	U-1895		
		9579359	Feb 10, 2029	U-1978		
<u>DEGARELIX ACETATE - FIRMAGON</u>						
N 022201	002	10695398	Apr 27, 2032	U-1895		
		10729739	Feb 10, 2029	U-1978		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>DEGARELIX ACETATE - FIRMAGON</u>						
N 022201	002	10973870	Feb 10, 2029	U-1978		
		11766468	Apr 27, 2032	U-1978		
		11826397	Apr 27, 2032	U-1978		
		9415085	Apr 27, 2032	U-1895		
		9579359	Feb 10, 2029	U-1978		
<u>DELAFLOXACIN MEGLUMINE - BAXDELA</u>						
N 208610	001	7728143	Jun 19, 2031	DS	NCE	Jun 19, 2022
		8252813	Oct 02, 2026	DP U-2028	GAIN	Jun 19, 2027
		8273892	Aug 06, 2026	DS		
		8648093	Oct 07, 2025	DP U-2028		
		8871938	Sep 23, 2029	DS		
		8969569	Oct 07, 2025	DP U-2028		
		9539250	Oct 07, 2025	DS DP U-2028		
		RE46617	Dec 28, 2029	DS		
<u>DELAFLOXACIN MEGLUMINE - BAXDELA</u>						
N 208611	001	12036219	Jun 02, 2034	U-3967	NCE	Jun 19, 2022
		12138257	May 01, 2032	DP	GAIN	Jun 19, 2027
		7635773	Mar 13, 2029	DP		
		7728143	Jun 19, 2031	DS		
		8252813	Oct 02, 2026	DP U-2028		
		8273892	Aug 06, 2026	DS		
		8410077	Mar 13, 2029	DP		
		8648093	Oct 07, 2025	DP U-2028		
		8871938	Sep 23, 2029	DS		
		9200088	Mar 13, 2029	DP		
		9493582	Feb 27, 2033	DP		
		9539250	Oct 07, 2025	DS DP U-2028		
		9750822	Mar 13, 2029	DP		
		RE46617	Dec 28, 2029	DS		
<u>DEOXYCHOLIC ACID - KYBELLA</u>						
N 206333	001	10500214	Mar 02, 2030	DP		
		7622130	Dec 10, 2027	U-1690		
		7754230	Dec 10, 2027	U-1690		
		8101593	Mar 02, 2030	DP		
		8242294	May 16, 2028	DS		
		8298556	Aug 03, 2025	U-1690		
		8367649	Mar 02, 2030	DP		
		8461140	Feb 21, 2028	DP		
		8546367	Feb 21, 2028	DP U-1690		
		8653058	Mar 02, 2030	DP		
		8846066	Feb 08, 2025	U-1690		
		8883770	Feb 21, 2028	DP		
		9522155	Feb 21, 2028	DP U-1940		
		9636349	Feb 21, 2028	U-1940		
		9949986	Feb 21, 2028	U-1940		
<u>DESMOPRESSIN ACETATE - NOCDURNA</u>						
N 022517	001	11020448	May 21, 2029	U-2327		
		11963995	May 21, 2029	U-2327		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>DESMOPRESSIN ACETATE - NOCDURNA</u>						
N 022517	001 9974826	Apr 13, 2030	U-2326			
<u>DESMOPRESSIN ACETATE - NOCDURNA</u>						
N 022517	002 10137167	May 21, 2029	U-2327			
	11963995	May 21, 2029	U-2327			
	9974826	Apr 13, 2030	U-2327			
<u>DESMOPRESSIN ACETATE - NOCTIVA</u>						
N 201656	001 11419914	Jun 15, 2030	U-3431			
	12090190	Jun 15, 2030	DP U-4003			
	9539302	Jun 15, 2030	DP			
<u>DESMOPRESSIN ACETATE - NOCTIVA</u>						
N 201656	002 11419914	Jun 15, 2030	U-3431			
	12090190	Jun 15, 2030	DP U-4003			
	9539302	Jun 15, 2030	DP			
<u>DESONIDE - VERDESO</u>						
N 021978	001 8460641	Aug 13, 2027	DP U-1412			
	8962000	Aug 31, 2025	DP U-1412			
	9492384	Aug 31, 2025	DP U-1412			
<u>DESOXIMETASONE - TOPICORT</u>						
N 204141	001 8277780	Sep 01, 2028	DP U-1408			
	8715624	May 26, 2026	DP U-1408			
<u>DESVENLAFAXINE SUCCINATE - PRISTIQ</u>						
N 021992	001 8269040	Jul 05, 2027	DS			
<u>DESVENLAFAXINE SUCCINATE - PRISTIQ</u>						
N 021992	002 8269040	Jul 05, 2027	DS			
<u>DESVENLAFAXINE SUCCINATE - PRISTIQ</u>						
N 021992	003 8269040	Jul 05, 2027	DS			
<u>DEUCRAVACITINIB - SOTYKTU</u>						
N 214958	001 10000480	Nov 07, 2033	DS DP		NCE	Sep 09, 2027
	11021475	Nov 07, 2033	U-3434			
	RE47929	Nov 07, 2033	DS DP U-3434			
<u>DEURUXOLITINIB PHOSPHATE - LEQSELVI</u>						
N 217900	001 10561659	May 04, 2037	U-3976		NCE	Jul 25, 2029
	11919907	May 21, 2041	DP U-3976			
	12076323	May 04, 2037	U-3976			
<u>DEUTETRABENAZINE - AUSTEDO</u>						
N 208082	001 10959996	Mar 07, 2036	U-3055			
	10959996*PED	Sep 07, 2036				
	11179386	Mar 15, 2038	DP U-1995			
	11179386	Mar 15, 2038	DP U-3055			
	11179386*PED	Sep 15, 2038				
	11357772	Mar 07, 2036	U-1995			
	11357772	Mar 07, 2036	U-3055			
	11357772*PED	Sep 07, 2036				
	11446291	Mar 07, 2036	U-1995			
	11446291*PED	Sep 07, 2036				

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>DEUTETRABENAZINE - AUSTEDO</u>						
N 208082 001	11564917	Mar 07, 2036	U-1995			
	11564917*PED	Sep 07, 2036				
	11648244	Mar 07, 2036	U-1995			
	11648244*PED	Sep 07, 2036				
	11666566	Sep 18, 2033	DP			
	11666566*PED	Mar 18, 2034				
	11813232	Mar 15, 2038	DP			
	11813232*PED	Sep 15, 2038				
	12016858	Mar 07, 2036	U-1995			
	12016858	Mar 07, 2036	U-3055			
	12016858*PED	Sep 07, 2036				
	8524733	Apr 03, 2031	DS DP			
	8524733*PED	Oct 03, 2031				
	9233959	Sep 18, 2033	DP			
	9233959*PED	Mar 18, 2034				
	9296739	Sep 18, 2033	DP			
	9296739*PED	Mar 18, 2034				
	9550780	Sep 18, 2033	DS DP U-1995			
	9550780	Sep 18, 2033	DS DP U-3055			
	9550780*PED	Mar 18, 2034				
	9814708	Sep 18, 2033	DP			
	9814708*PED	Mar 18, 2034				
<u>DEUTETRABENAZINE - AUSTEDO</u>						
N 208082 002	10959996	Mar 07, 2036	U-3055			
	10959996*PED	Sep 07, 2036				
	11179386	Mar 15, 2038	DP U-1995			
	11179386	Mar 15, 2038	DP U-3055			
	11179386*PED	Sep 15, 2038				
	11357772	Mar 07, 2036	U-1995			
	11357772	Mar 07, 2036	U-3055			
	11357772*PED	Sep 07, 2036				
	11446291	Mar 07, 2036	U-1995			
	11446291*PED	Sep 07, 2036				
	11564917	Mar 07, 2036	U-1995			
	11564917*PED	Sep 07, 2036				
	11648244	Mar 07, 2036	U-1995			
	11648244*PED	Sep 07, 2036				
	11666566	Sep 18, 2033	DP			
	11666566*PED	Mar 18, 2034				
	11813232	Mar 15, 2038	DP			
	11813232*PED	Sep 15, 2038				
	12016858	Mar 07, 2036	U-1995			
	12016858	Mar 07, 2036	U-3055			
	12016858*PED	Sep 07, 2036				
	8524733	Apr 03, 2031	DS DP			
	8524733*PED	Oct 03, 2031				
	9233959	Sep 18, 2033	DP			
	9233959*PED	Mar 18, 2034				
	9296739	Sep 18, 2033	DP			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>DEUTETRABENAZINE - AUSTEDO</u>						
N 208082 002	9296739*PED	Mar 18, 2034				
	9550780	Sep 18, 2033	DS DP U-1995			
	9550780	Sep 18, 2033	DS DP U-3055			
	9550780*PED	Mar 18, 2034				
	9814708	Sep 18, 2033	DP			
	9814708*PED	Mar 18, 2034				
<u>DEUTETRABENAZINE - AUSTEDO</u>						
N 208082 003	10959996	Mar 07, 2036	U-3055			
	10959996*PED	Sep 07, 2036				
	11179386	Mar 15, 2038	DP U-1995			
	11179386	Mar 15, 2038	DP U-3055			
	11179386*PED	Sep 15, 2038				
	11357772	Mar 07, 2036	U-1995			
	11357772	Mar 07, 2036	U-3055			
	11357772*PED	Sep 07, 2036				
	11446291	Mar 07, 2036	U-1995			
	11446291*PED	Sep 07, 2036				
	11564917	Mar 07, 2036	U-1995			
	11564917*PED	Sep 07, 2036				
	11648244	Mar 07, 2036	U-1995			
	11648244*PED	Sep 07, 2036				
	11666566	Sep 18, 2033	DP			
	11666566*PED	Mar 18, 2034				
	11813232	Mar 15, 2038	DP			
	11813232*PED	Sep 15, 2038				
	12016858	Mar 07, 2036	U-1995			
	12016858	Mar 07, 2036	U-3055			
	12016858*PED	Sep 07, 2036				
	8524733	Apr 03, 2031	DS DP			
	8524733*PED	Oct 03, 2031				
	9233959	Sep 18, 2033	DP			
	9233959*PED	Mar 18, 2034				
	9296739	Sep 18, 2033	DP			
	9296739*PED	Mar 18, 2034				
	9550780	Sep 18, 2033	DS DP U-1995			
	9550780	Sep 18, 2033	DS DP U-3055			
	9550780*PED	Mar 18, 2034				
	9814708	Sep 18, 2033	DP			
	9814708*PED	Mar 18, 2034				
<u>DEUTETRABENAZINE - AUSTEDO XR</u>						
N 216354 001	10959996	Mar 07, 2036	U-3055			
	10959996*PED	Sep 07, 2036				
	11179386	Mar 15, 2038	DP U-1995			
	11179386	Mar 15, 2038	DP U-3055			
	11179386*PED	Sep 15, 2038				
	11311488	Jun 10, 2041	DP U-1995			
	11311488	Jun 10, 2041	DP U-3055			
	11357772	Mar 07, 2036	U-1995			
	11357772	Mar 07, 2036	U-3055			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>DEUTETRABENAZINE - AUSTEDO XR</u>						
N 216354 001	11357772*PED	Sep 07, 2036				
	11446291	Mar 07, 2036	U-1995			
	11446291*PED	Sep 07, 2036				
	11564917	Mar 07, 2036	U-1995			
	11564917*PED	Sep 07, 2036				
	11648244	Mar 07, 2036	U-1995			
	11648244*PED	Sep 07, 2036				
	11813232	Mar 15, 2038	DP			
	11813232*PED	Sep 15, 2038				
	12016858	Mar 07, 2036	U-1995			
	12016858	Mar 07, 2036	U-3055			
	12016858*PED	Sep 07, 2036				
	8524733	Apr 03, 2031	DS DP			
	8524733*PED	Oct 03, 2031				
	9550780	Sep 18, 2033	DS DP U-1995			
	9550780	Sep 18, 2033	DS DP U-3055			
	9550780*PED	Mar 18, 2034				
<u>DEUTETRABENAZINE - AUSTEDO XR</u>						
N 216354 002	10959996	Mar 07, 2036	U-3055			
	10959996*PED	Sep 07, 2036				
	11179386	Mar 15, 2038	DP U-1995			
	11179386	Mar 15, 2038	DP U-3055			
	11179386*PED	Sep 15, 2038				
	11311488	Jun 10, 2041	DP U-1995			
	11311488	Jun 10, 2041	DP U-3055			
	11357772	Mar 07, 2036	U-1995			
	11357772	Mar 07, 2036	U-3055			
	11357772*PED	Sep 07, 2036				
	11446291	Mar 07, 2036	U-1995			
	11446291*PED	Sep 07, 2036				
	11564917	Mar 07, 2036	U-1995			
	11564917*PED	Sep 07, 2036				
	11648244	Mar 07, 2036	U-1995			
	11648244*PED	Sep 07, 2036				
	11813232	Mar 15, 2038	DP			
	11813232*PED	Sep 15, 2038				
	12016858	Mar 07, 2036	U-1995			
	12016858	Mar 07, 2036	U-3055			
	12016858*PED	Sep 07, 2036				
	8524733	Apr 03, 2031	DS DP			
	8524733*PED	Oct 03, 2031				
	9550780	Sep 18, 2033	DS DP U-1995			
	9550780	Sep 18, 2033	DS DP U-3055			
	9550780*PED	Mar 18, 2034				
<u>DEUTETRABENAZINE - AUSTEDO XR</u>						
N 216354 003	10959996	Mar 07, 2036	U-3055			
	10959996*PED	Sep 07, 2036				
	11179386	Mar 15, 2038	DP U-1995			
	11179386	Mar 15, 2038	DP U-3055			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>DEUTETRABENAZINE - AUSTEDO XR</u>						
N 216354 003	11179386*PED	Sep 15, 2038				
	11311488	Jun 10, 2041	DP U-1995			
	11311488	Jun 10, 2041	DP U-3055			
	11357772	Mar 07, 2036	U-1995			
	11357772	Mar 07, 2036	U-3055			
	11357772*PED	Sep 07, 2036				
	11446291	Mar 07, 2036	U-1995			
	11446291*PED	Sep 07, 2036				
	11564917	Mar 07, 2036	U-1995			
	11564917*PED	Sep 07, 2036				
	11648244	Mar 07, 2036	U-1995			
	11648244*PED	Sep 07, 2036				
	11813232	Mar 15, 2038	DP			
	11813232*PED	Sep 15, 2038				
	12016858	Mar 07, 2036	U-1995			
	12016858	Mar 07, 2036	U-3055			
	12016858*PED	Sep 07, 2036				
	8524733	Apr 03, 2031	DS DP			
	8524733*PED	Oct 03, 2031				
	9550780	Sep 18, 2033	DS DP U-1995			
	9550780	Sep 18, 2033	DS DP U-3055			
	9550780*PED	Mar 18, 2034				
<u>DEUTETRABENAZINE - AUSTEDO XR</u>						
N 216354 004	10959996	Mar 07, 2036	U-3055			
	10959996*PED	Sep 07, 2036				
	11179386	Mar 15, 2038	DP U-1995			
	11179386	Mar 15, 2038	DP U-3055			
	11179386*PED	Sep 15, 2038				
	11311488	Jun 10, 2041	DP U-1995			
	11311488	Jun 10, 2041	DP U-3055			
	11357772	Mar 07, 2036	U-1995			
	11357772	Mar 07, 2036	U-3055			
	11357772*PED	Sep 07, 2036				
	11446291	Mar 07, 2036	U-1995			
	11446291*PED	Sep 07, 2036				
	11564917	Mar 07, 2036	U-1995			
	11564917*PED	Sep 07, 2036				
	11648244	Mar 07, 2036	U-1995			
	11648244*PED	Sep 07, 2036				
	11813232	Mar 15, 2038	DP			
	11813232*PED	Sep 15, 2038				
	12016858	Mar 07, 2036	U-1995			
	12016858	Mar 07, 2036	U-3055			
	12016858*PED	Sep 07, 2036				
	8524733	Apr 03, 2031	DS DP			
	8524733*PED	Oct 03, 2031				
	9550780	Sep 18, 2033	DS DP U-1995			
	9550780	Sep 18, 2033	DS DP U-3055			
	9550780*PED	Mar 18, 2034				

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>DEUTETRABENAZINE - AUSTEDO XR</u>						
N 216354 004	10959996	Mar 07, 2036	U-3055			
	10959996*PED	Sep 07, 2036				
	11179386	Mar 15, 2038	DP U-1995			
	11179386	Mar 15, 2038	DP U-3055			
	11179386*PED	Sep 15, 2038				
	11311488	Jun 10, 2041	DP U-1995			
	11311488	Jun 10, 2041	DP U-3055			
	11357772	Mar 07, 2036	U-1995			
	11357772	Mar 07, 2036	U-3055			
	11357772*PED	Sep 07, 2036				
	11446291	Mar 07, 2036	U-1995			
	11446291*PED	Sep 07, 2036				
	11564917	Mar 07, 2036	U-1995			
	11564917*PED	Sep 07, 2036				
	11648244	Mar 07, 2036	U-1995			
	11648244*PED	Sep 07, 2036				
	11813232	Mar 15, 2038	DP			
	11813232*PED	Sep 15, 2038				
	12016858	Mar 07, 2036	U-1995			
	12016858	Mar 07, 2036	U-3055			
	12016858*PED	Sep 07, 2036				
	8524733	Apr 03, 2031	DS DP			
	8524733*PED	Oct 03, 2031				
	9550780	Sep 18, 2033	DS DP U-1995			
	9550780	Sep 18, 2033	DS DP U-3055			
	9550780*PED	Mar 18, 2034				
<u>DEUTETRABENAZINE - AUSTEDO XR</u>						
N 216354 005	10959996	Mar 07, 2036	U-3055			
	10959996*PED	Sep 07, 2036				
	11179386	Mar 15, 2038	DP U-1995			
	11179386	Mar 15, 2038	DP U-3055			
	11179386*PED	Sep 15, 2038				
	11311488	Jun 10, 2041	DP U-1995			
	11311488	Jun 10, 2041	DP U-3055			
	11357772	Mar 07, 2036	U-1995			
	11357772	Mar 07, 2036	U-3055			
	11357772*PED	Sep 07, 2036				
	11446291	Mar 07, 2036	U-1995			
	11446291*PED	Sep 07, 2036				
	11564917	Mar 07, 2036	U-1995			
	11564917*PED	Sep 07, 2036				
	11648244	Mar 07, 2036	U-1995			
	11648244*PED	Sep 07, 2036				
	11813232	Mar 15, 2038	DP			
	11813232*PED	Sep 15, 2038				
	12016858	Mar 07, 2036	U-1995			
	12016858	Mar 07, 2036	U-3055			
	12016858*PED	Sep 07, 2036				
	8524733	Apr 03, 2031	DS DP			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>DEUTETRABENAZINE - AUSTEDO XR</u>						
N 216354 005	8524733*PED	Oct 03, 2031				
	9550780	Sep 18, 2033	DS DP	U-1995		
	9550780	Sep 18, 2033	DS DP	U-3055		
	9550780*PED	Mar 18, 2034				
<u>DEUTETRABENAZINE - AUSTEDO XR</u>						
N 216354 006	10959996	Mar 07, 2036		U-3055		
	10959996*PED	Sep 07, 2036				
	11179386	Mar 15, 2038	DP	U-1995		
	11179386	Mar 15, 2038	DP	U-3055		
	11179386*PED	Sep 15, 2038				
	11311488	Jun 10, 2041	DP	U-1995		
	11311488	Jun 10, 2041	DP	U-3055		
	11357772	Mar 07, 2036		U-1995		
	11357772	Mar 07, 2036		U-3055		
	11357772*PED	Sep 07, 2036				
	11446291	Mar 07, 2036		U-1995		
	11446291*PED	Sep 07, 2036				
	11564917	Mar 07, 2036		U-1995		
	11564917*PED	Sep 07, 2036				
	11648244	Mar 07, 2036		U-1995		
	11648244*PED	Sep 07, 2036				
	11813232	Mar 15, 2038	DP			
	11813232*PED	Sep 15, 2038				
	12016858	Mar 07, 2036		U-1995		
	12016858	Mar 07, 2036		U-3055		
	12016858*PED	Sep 07, 2036				
	8524733	Apr 03, 2031	DS DP			
	8524733*PED	Oct 03, 2031				
	9550780	Sep 18, 2033	DS DP	U-1995		
	9550780	Sep 18, 2033	DS DP	U-3055		
	9550780*PED	Mar 18, 2034				
<u>DEUTETRABENAZINE - AUSTEDO XR</u>						
N 216354 007	10959996	Mar 07, 2036		U-3055		
	10959996*PED	Sep 07, 2036				
	11179386	Mar 15, 2038	DP	U-1995		
	11179386	Mar 15, 2038	DP	U-3055		
	11179386*PED	Sep 15, 2038				
	11311488	Jun 10, 2041	DP	U-1995		
	11311488	Jun 10, 2041	DP	U-3055		
	11357772	Mar 07, 2036		U-1995		
	11357772	Mar 07, 2036		U-3055		
	11357772*PED	Sep 07, 2036				
	11446291	Mar 07, 2036		U-1995		
	11446291*PED	Sep 07, 2036				
	11564917	Mar 07, 2036		U-1995		
	11564917*PED	Sep 07, 2036				
	11648244	Mar 07, 2036		U-1995		
	11648244*PED	Sep 07, 2036				
	11813232	Mar 15, 2038	DP			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>DEUTETRABENAZINE - AUSTEDO XR</u>						
N 216354 007	11813232*PED	Sep 15, 2038				
	12016858	Mar 07, 2036	U-1995			
	12016858	Mar 07, 2036	U-3055			
	12016858*PED	Sep 07, 2036				
	8524733	Apr 03, 2031	DS DP			
	8524733*PED	Oct 03, 2031				
	9550780	Sep 18, 2033	DS DP U-1995			
	9550780	Sep 18, 2033	DS DP U-3055			
	9550780*PED	Mar 18, 2034				
<u>DEUTETRABENAZINE - AUSTEDO XR</u>						
N 216354 008	10959996	Mar 07, 2036	U-3055			
	10959996*PED	Sep 07, 2036				
	11179386	Mar 15, 2038	DP U-1995			
	11179386	Mar 15, 2038	DP U-3055			
	11179386*PED	Sep 15, 2038				
	11311488	Jun 10, 2041	DP U-1995			
	11311488	Jun 10, 2041	DP U-3055			
	11357772	Mar 07, 2036	U-1995			
	11357772	Mar 07, 2036	U-3055			
	11357772*PED	Sep 07, 2036				
	11446291	Mar 07, 2036	U-1995			
	11446291*PED	Sep 07, 2036				
	11564917	Mar 07, 2036	U-1995			
	11564917*PED	Sep 07, 2036				
	11648244	Mar 07, 2036	U-1995			
	11648244*PED	Sep 07, 2036				
	11813232	Mar 15, 2038	DP			
	11813232*PED	Sep 15, 2038				
	12016858	Mar 07, 2036	U-1995			
	12016858	Mar 07, 2036	U-3055			
	12016858*PED	Sep 07, 2036				
	8524733	Apr 03, 2031	DS DP			
	8524733*PED	Oct 03, 2031				
	9550780	Sep 18, 2033	DS DP U-1995			
	9550780	Sep 18, 2033	DS DP U-3055			
	9550780*PED	Mar 18, 2034				
<u>DEXAMETHASONE - DEXTENZA</u>						
N 208742 001	11458041	Nov 16, 2037	U-1680			
	11458041	Nov 16, 2037	U-3455			
	12144889	Apr 26, 2041	U-1680			
	12150896	Oct 07, 2036	U-1680			
	12150896	Oct 07, 2036	U-3455			
	8409606	May 14, 2030	DP			
	8563027	Feb 12, 2030	U-2487			
<u>DEXAMETHASONE - DEXYCU KIT</u>						
N 208912 001	10022502	Jun 22, 2034	U-2340			
	10028965	May 23, 2034	U-2340			
	10159683	May 23, 2034	DP			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>DEXAMETHASONE - DEXYCU KIT</u>						
N 208912	001 10799642	May 11, 2032	DP			
	11097061	Jun 23, 2039	DP U-3418			
<u>DEXAMETHASONE - HEMADY</u>						
N 211379	001 10537585	Dec 18, 2037	DP		ODE-271	Oct 03, 2026
	11304961	Dec 18, 2037	DP U-3344			
<u>DEXAMETHASONE; TOBRAMYCIN - TOBRADEX ST</u>						
N 050818	001 7795316	Aug 03, 2028	DP U-1082			
	8101582	Dec 19, 2027	DP U-1082			
	8450287	Dec 19, 2027	DP			
<u>DEXLANSOPRAZOLE - DEXILANT</u>						
N 022287	001 7790755	Aug 02, 2026	DP			
	8105626	Sep 27, 2026	DP			
	8173158	Mar 17, 2030	U-949			
	8173158	Mar 17, 2030	U-950			
	8173158	Mar 17, 2030	U-951			
	8461187	Jan 17, 2026	DP			
	8461187*PED	Jul 17, 2026				
	8871273	Jan 11, 2028	DP			
	9011926	Feb 24, 2026	DP			
	9233103	Mar 05, 2032	U-1805			
	9238029	Jan 17, 2026	DP			
<u>DEXLANSOPRAZOLE - DEXILANT</u>						
N 022287	002 7790755	Aug 02, 2026	DP			
	8105626	Sep 27, 2026	DP			
	8173158	Mar 17, 2030	U-949			
	8173158	Mar 17, 2030	U-950			
	8173158	Mar 17, 2030	U-951			
	8461187	Jan 17, 2026	DP			
	8461187*PED	Jul 17, 2026				
	8871273	Jan 11, 2028	DP			
	9011926	Feb 24, 2026	DP			
	9233103	Mar 05, 2032	U-1805			
	9238029	Jan 17, 2026	DP			
<u>DEXLANSOPRAZOLE - DEXILANT SOLUTAB</u>						
N 208056	001 8461187	Jan 17, 2026	DP			
	8461187*PED	Jul 17, 2026				
	8871273	Jan 11, 2028	DP			
	8871273*PED	Jul 11, 2028				
	9011926	Feb 24, 2026	DP			
	9238029	Jan 17, 2026	DP			
	9241910	Mar 10, 2029	DP			
<u>DEXMEDETOMIDINE HYDROCHLORIDE - DEXMEDETOMIDINE HYDROCHLORIDE</u>						
A 218112	001				CGT	Mar 30, 2025
<u>DEXMEDETOMIDINE HYDROCHLORIDE - DEXMEDETOMIDINE HYDROCHLORIDE</u>						
A 218112	002				CGT	Mar 30, 2025

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>DEXMEDETOMIDINE HYDROCHLORIDE - PRECEDEX</u>						
N 021038	001				NPP	Dec 16, 2025
<u>DEXMEDETOMIDINE HYDROCHLORIDE - PRECEDEX</u>						
N 021038	002	10016396	Jan 04, 2032	DP	NPP	Dec 16, 2025
		8242158	Jan 04, 2032	DP		
		8242158*PED	Jul 04, 2032			
		8338470	Jan 04, 2032	DP		
		8338470*PED	Jul 04, 2032			
		8455527	Jan 04, 2032	U-421		
		8455527*PED	Jul 04, 2032			
		8648106	Jan 04, 2032	DP		
		8648106*PED	Jul 04, 2032			
		9320712	Jan 04, 2032	DP		
		9320712*PED	Jul 04, 2032			
		9616049	Jan 04, 2032	DP		
		9616049*PED	Jul 04, 2032			
<u>DEXMEDETOMIDINE HYDROCHLORIDE - PRECEDEX</u>						
N 021038	003	10016396	Jan 04, 2032	DP	NPP	Dec 16, 2025
		8242158	Jan 04, 2032	DP		
		8242158*PED	Jul 04, 2032			
		8338470	Jan 04, 2032	DP		
		8338470*PED	Jul 04, 2032			
		8455527	Jan 04, 2032	U-421		
		8455527*PED	Jul 04, 2032			
		8648106	Jan 04, 2032	DP		
		8648106*PED	Jul 04, 2032			
		9320712	Jan 04, 2032	DP		
		9320712*PED	Jul 04, 2032			
		9616049	Jan 04, 2032	DP		
		9616049*PED	Jul 04, 2032			
<u>DEXMEDETOMIDINE HYDROCHLORIDE - PRECEDEX</u>						
N 021038	004	8242158	Jan 04, 2032	DP	NPP	Dec 16, 2025
		8242158*PED	Jul 04, 2032			
		8338470	Jan 04, 2032	DP		
		8338470*PED	Jul 04, 2032			
		8455527	Jan 04, 2032	U-421		
		8455527*PED	Jul 04, 2032			
		8648106	Jan 04, 2032	DP		
		8648106*PED	Jul 04, 2032			
		9320712	Jan 04, 2032	DP		
		9320712*PED	Jul 04, 2032			
		9616049	Jan 04, 2032	DP		
		9616049*PED	Jul 04, 2032			
<u>DEXMEDETOMIDINE HYDROCHLORIDE - PRECEDEX</u>						
N 021038	005				NPP	Dec 16, 2025
<u>DEXMEDETOMIDINE HYDROCHLORIDE - DEXMEDETOMIDINE HYDROCHLORIDE</u>						
N 206628	003	9649296	Apr 20, 2036	DP		
		9717796	Apr 20, 2036	DP		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>DEXMEDETOMIDINE HYDROCHLORIDE - DEXMEDETOMIDINE HYDROCHLORIDE</u>						
N 206628	004	9649296	Apr 20, 2036	DP		
		9717796	Apr 20, 2036	DP		
<u>DEXMEDETOMIDINE HYDROCHLORIDE - IGALMI</u>						
N 215390	001	10792246	Jun 26, 2039	DP U-3350	NP	Apr 05, 2025
		11478422	Jun 26, 2039	DP		
		11497711	Jun 26, 2039	DP U-3645		
		11517524	Jun 26, 2039	DP U-3645		
		11786508	Dec 29, 2037	U-3698		
		11806334	Jan 12, 2043	U-3725		
		11839604	Dec 29, 2037	U-3756		
		11890272	Jul 17, 2040	U-3756		
		11890272	Jul 17, 2040	U-3935		
		11998528	Jan 12, 2043	U-3935		
		11998529	Jul 17, 2040	U-3935		
		12090140	Jan 12, 2043	U-3986		
		12109196	Jul 17, 2040	U-3935		
		12138247	Jan 12, 2043	U-4029		
<u>DEXMEDETOMIDINE HYDROCHLORIDE - IGALMI</u>						
N 215390	002	10792246	Jun 26, 2039	DP U-3350	NP	Apr 05, 2025
		11478422	Jun 26, 2039	DP		
		11497711	Jun 26, 2039	DP U-3645		
		11517524	Jun 26, 2039	DP U-3645		
		11786508	Dec 29, 2037	U-3698		
		11806334	Jan 12, 2043	U-3725		
		11839604	Dec 29, 2037	U-3756		
		11890272	Jul 17, 2040	U-3756		
		11890272	Jul 17, 2040	U-3935		
		11998528	Jan 12, 2043	U-3935		
		11998529	Jul 17, 2040	U-3935		
		12090140	Jan 12, 2043	U-3986		
		12109196	Jul 17, 2040	U-3935		
		12138247	Jan 12, 2043	U-4029		
<u>DEXMETHYLPHENIDATE HYDROCHLORIDE; SERDEXMETHYLPHENIDATE CHLORIDE - AZSTARYS</u>						
N 212994	001	10584112	Dec 09, 2037	DS DP	NCE	May 07, 2026
		10584113	Dec 09, 2037	DP		
		10759778	Dec 09, 2037	DP		
		10858341	Dec 09, 2037	U-3094		
		10954213	Dec 09, 2037	U-3094		
		9079928	Jul 27, 2032	DP		
<u>DEXMETHYLPHENIDATE HYDROCHLORIDE; SERDEXMETHYLPHENIDATE CHLORIDE - AZSTARYS</u>						
N 212994	002	10584112	Dec 09, 2037	DS DP	NCE	May 07, 2026
		10584113	Dec 09, 2037	DP		
		10759778	Dec 09, 2037	DP		
		10858341	Dec 09, 2037	U-3094		
		10954213	Dec 09, 2037	U-3094		
		9079928	Jul 27, 2032	DP		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>DEXMETHYLPHENIDATE HYDROCHLORIDE; SERDEXMETHYLPHENIDATE CHLORIDE - AZSTARYS</u>						
N 212994 003	10584112	Dec 09, 2037	DS DP		NCE	May 07, 2026
	10584113	Dec 09, 2037	DP			
	10759778	Dec 09, 2037	DP			
	10858341	Dec 09, 2037	U-3094			
	10954213	Dec 09, 2037	U-3094			
	9079928	Jul 27, 2032	DP			
<u>DEXTROAMPHETAMINE - XELSTRYM</u>						
N 215401 001	11559501	Jan 06, 2042	DP		NP	Mar 22, 2025
	8591941	Oct 07, 2025	DP			
	8632802	Oct 07, 2025	DP			
	9034370	Oct 07, 2025	DP			
	9456993	Oct 24, 2033	DP U-3340			
	9474722	Oct 24, 2033	DP			
<u>DEXTROAMPHETAMINE - XELSTRYM</u>						
N 215401 002	11559501	Jan 06, 2042	DP		NP	Mar 22, 2025
	8591941	Oct 07, 2025	DP			
	8632802	Oct 07, 2025	DP			
	9034370	Oct 07, 2025	DP			
	9456993	Oct 24, 2033	DP U-3340			
	9474722	Oct 24, 2033	DP			
<u>DEXTROAMPHETAMINE - XELSTRYM</u>						
N 215401 003	11559501	Jan 06, 2042	DP		NP	Mar 22, 2025
	8591941	Oct 07, 2025	DP			
	8632802	Oct 07, 2025	DP			
	9034370	Oct 07, 2025	DP			
	9456993	Oct 24, 2033	DP U-3340			
	9474722	Oct 24, 2033	DP			
<u>DEXTROAMPHETAMINE - XELSTRYM</u>						
N 215401 004	11559501	Jan 06, 2042	DP		NP	Mar 22, 2025
	8591941	Oct 07, 2025	DP			
	8632802	Oct 07, 2025	DP			
	9034370	Oct 07, 2025	DP			
	9456993	Oct 24, 2033	DP U-3340			
	9474722	Oct 24, 2033	DP			
<u>DEXTROMETHORPHAN HYDROBROMIDE; QUINIDINE SULFATE - NUEDEXTA</u>						
N 021879 001	7659282	Aug 13, 2026	U-1093			
<u>DIAZEPAM - VALTOCO</u>						
N 211635 001	10265402	May 11, 2025	DP		ODE-279	Jan 10, 2027
	11241414	Mar 27, 2029	DP			
	11793786	Mar 27, 2029	DP			
	8895546	Mar 27, 2029	DP			
	8927497	Jul 21, 2025	DP U-2727			
	9642913	May 11, 2025	DP			
<u>DIAZEPAM - VALTOCO</u>						
N 211635 002	10265402	May 11, 2025	DP		ODE-279	Jan 10, 2027
	11241414	Mar 27, 2029	DP			
	11793786	Mar 27, 2029	DP			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>DIAZEPAM - VALTOCO</u>						
N 211635	002	8895546	Mar 27, 2029	DP		
		8927497	Jul 21, 2025	DP	U-2727	
		9642913	May 11, 2025	DP		
<u>DIAZEPAM - VALTOCO</u>						
N 211635	003	10265402	May 11, 2025	DP	ODE-279	Jan 10, 2027
		11241414	Mar 27, 2029	DP		
		11793786	Mar 27, 2029	DP		
		8895546	Mar 27, 2029	DP		
		8927497	Jul 21, 2025	DP	U-2727	
		9642913	May 11, 2025	DP		
<u>DIAZEPAM - LIBERVANT</u>						
N 218623	001	11273131	Jun 18, 2038	DP	NP	Apr 26, 2027
					ODE-501	Apr 26, 2031
<u>DIAZEPAM - LIBERVANT</u>						
N 218623	002	11273131	Jun 18, 2038	DP	NP	Apr 26, 2027
					ODE-501	Apr 26, 2031
<u>DIAZEPAM - LIBERVANT</u>						
N 218623	003	11273131	Jun 18, 2038	DP	NP	Apr 26, 2027
					ODE-501	Apr 26, 2031
<u>DIAZEPAM - LIBERVANT</u>						
N 218623	004	11273131	Jun 18, 2038	DP	NP	Apr 26, 2027
					ODE-501	Apr 26, 2031
<u>DIAZEPAM - LIBERVANT</u>						
N 218623	005	11273131	Jun 18, 2038	DP	NP	Apr 26, 2027
					ODE-501	Apr 26, 2031
<u>DICLOFENAC - ZORVOLEX</u>						
N 204592	001	8679544	Apr 23, 2030	DP		
		8999387	Apr 23, 2030		U-55	
		9017721	Apr 23, 2030	DP		
		9173854	Apr 23, 2030	DP		
		9180095	Apr 23, 2030		U-55	
		9180096	Apr 23, 2030	DP		
		9186328	Apr 23, 2030		U-55	
<u>DICLOFENAC - ZORVOLEX</u>						
N 204592	002	8679544	Apr 23, 2030	DP		
		8999387	Apr 23, 2030		U-55	
		9017721	Apr 23, 2030	DP		
		9173854	Apr 23, 2030	DP		
		9180095	Apr 23, 2030		U-55	
		9180096	Apr 23, 2030	DP		
		9186328	Apr 23, 2030		U-55	
<u>DICLOFENAC EPOLAMINE - LICART</u>						
N 206976	001	11344520	Feb 20, 2035		U-3393	
		11351133	Feb 20, 2035		U-3393	

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>DICLOFENAC POTASSIUM - CAMBIA</u>						
N 022165	001 7759394	Jun 16, 2026	DS DP U-436			
	8097651	Jun 16, 2026	DS DP U-436			
	8927604	Jun 16, 2026	U-436			
	9827197	Jun 16, 2026	DP			
<u>DICLOFENAC POTASSIUM - ZIPSOR</u>						
N 022202	001 7662858	Feb 24, 2029	U-1035			
	7884095	Feb 24, 2029	U-1111			
	7939518	Feb 24, 2029	U-980			
	8110606	Feb 24, 2029	U-980			
	8623920	Feb 24, 2029	U-1482			
	9561200	Feb 24, 2029	U-1482			
<u>DICLOFENAC SODIUM - PENNSAID</u>						
N 020947	001 8217078	Jul 10, 2029	U-1248			
	8546450	Aug 09, 2030	U-1435			
	8546450	Aug 09, 2030	U-1436			
	8618164	Jul 10, 2029	U-1477			
	8741956	Jul 10, 2029	U-1435			
<u>DICLOFENAC SODIUM - DYLOJECT</u>						
N 022396	001 8946292	Mar 22, 2027	U-1659			
<u>DICLOFENAC SODIUM - PENNSAID</u>						
N 204623	001 8217078	Jul 10, 2029	U-1477			
	8252838	Apr 21, 2028	DP U-1489			
	8546450	Aug 09, 2030	U-1435			
	8546450	Aug 09, 2030	U-1436			
	8563613	Oct 17, 2027	DP U-1488			
	8618164	Jul 10, 2029	U-1477			
	8741956	Jul 10, 2029	U-1435			
	8871809	Oct 17, 2027	U-1614			
	9066913	Oct 17, 2027	DP U-1488			
	9101591	Oct 17, 2027	DP U-1488			
	9132110	Oct 17, 2027	U-1488			
	9168304	Oct 17, 2027	DP			
	9168305	Oct 17, 2027	U-1488			
	9220784	Oct 17, 2027	U-1488			
	9339551	Oct 17, 2027	U-1488			
	9339552	Oct 17, 2027	DP U-1488			
	9370501	Jul 10, 2029	U-1614			
	9375412	Jul 10, 2029	U-1614			
	9415029	Jul 10, 2029	U-1614			
	9539335	Oct 17, 2027	U-1614			
<u>DIENOEST; DIENOEST; DIENOEST; ESTRADIOL VALERATE; ESTRADIOL VALERATE; ESTRADIOL VALERATE; ESTRADIOL VALERATE; ESTRADIOL VALERATE - NATAZIA</u>						
N 022252	001 8071577	May 13, 2026	DP U-1			
	8153616	Jan 30, 2028	U-1240			
<u>DIFELIKEFALIN ACETATE - KORSUVA</u>						
N 214916	001 10017536	Nov 12, 2027	DS U-3204		NCE	Aug 23, 2026
	10138270	Nov 12, 2027	U-3204			
	10793596	Nov 12, 2027	DS DP U-3204			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>DIFELIKEFALIN ACETATE - KORSUVA</u>						
N 214916	001 7402564	Nov 12, 2027	DS DP U-3204			
	7713937	Nov 12, 2027	DS DP U-3204			
	7727963	Nov 12, 2027	DS DP U-3204			
	8217007	Nov 12, 2027	U-3204			
	8236766	Nov 12, 2027	U-3204			
	8486894	Nov 12, 2027	U-3204			
	8536131	Nov 12, 2027	DS DP U-3204			
	9334305	Nov 12, 2027	U-3204			
	9359399	Nov 12, 2027	U-3204			
<u>DIHYDROERGOTAMINE MESYLATE - TRUDHESA</u>						
N 213436	001 10507295	Dec 25, 2032	DP			
	10940278	Jan 23, 2033	DP			
	11185497	Jan 04, 2039	U-3218			
	11266799	Nov 05, 2036	DP			
	9550036	Sep 05, 2032	DP			
	9919117	Mar 17, 2033	DP U-3218			
<u>DIMETHYL FUMARATE - TECFIDERA</u>						
N 204063	001 10391160	Mar 13, 2035	U-3148			
	10555993	Mar 13, 2035	U-3148			
	10959972	Nov 16, 2035	U-1384			
	10994003	Mar 13, 2035	U-3148			
	11007166	Nov 16, 2035	U-1384			
	11007167	Nov 16, 2035	U-1384			
	11129806	Nov 16, 2035	U-1384			
	11246850	Nov 16, 2035	U-1384			
	8399514	Feb 07, 2028	U-1384			
<u>DIMETHYL FUMARATE - TECFIDERA</u>						
N 204063	002 10391160	Mar 13, 2035	U-3148			
	10555993	Mar 13, 2035	U-3148			
	10959972	Nov 16, 2035	U-1384			
	10994003	Mar 13, 2035	U-3148			
	11007166	Nov 16, 2035	U-1384			
	11007167	Nov 16, 2035	U-1384			
	11129806	Nov 16, 2035	U-1384			
	11246850	Nov 16, 2035	U-1384			
	8399514	Feb 07, 2028	U-1384			
<u>DIROXIMEL FUMARATE - VUMERITY</u>						
N 211855	001 10080733	Sep 20, 2033	DS DP U-1384			
	8669281	Sep 20, 2033	DS DP			
	9090558	Sep 20, 2033	U-1384			
<u>DOCETAXEL - DOCETAXEL</u>						
N 205934	001 10842770	Aug 07, 2031	DP U-2998			
	8940786	Sep 30, 2033	DP U-1789			
	9308195	Sep 30, 2033	DP			
	9763880	Sep 30, 2033	U-2558			
	9763880	Sep 30, 2033	U-2559			
	9763880	Sep 30, 2033	U-2560			
	9763880	Sep 30, 2033	U-2561			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>DOCETAXEL - DOCETAXEL</u>						
N 205934	001	9763880	Sep 30, 2033	U-2562		
		9763880	Sep 30, 2033	U-2563		
		9763880	Sep 30, 2033	U-2564		
<u>DOCETAXEL - DOCETAXEL</u>						
N 205934	002	10842770	Aug 07, 2031	DP U-2998		
		8940786	Sep 30, 2033	DP U-1789		
		9308195	Sep 30, 2033	DP		
		9763880	Sep 30, 2033	U-2558		
		9763880	Sep 30, 2033	U-2559		
		9763880	Sep 30, 2033	U-2560		
		9763880	Sep 30, 2033	U-2561		
		9763880	Sep 30, 2033	U-2562		
		9763880	Sep 30, 2033	U-2563		
		9763880	Sep 30, 2033	U-2564		
<u>DOCETAXEL - DOCETAXEL</u>						
N 205934	003	10842770	Aug 07, 2031	DP U-2998		
		8940786	Sep 30, 2033	DP U-1789		
		9308195	Sep 30, 2033	DP		
		9763880	Sep 30, 2033	U-2558		
		9763880	Sep 30, 2033	U-2559		
		9763880	Sep 30, 2033	U-2560		
		9763880	Sep 30, 2033	U-2561		
		9763880	Sep 30, 2033	U-2562		
		9763880	Sep 30, 2033	U-2563		
		9763880	Sep 30, 2033	U-2564		
<u>DOCETAXEL - DOCIVYX</u>						
N 215813	001	10398785	Mar 14, 2036	DP		
<u>DOCETAXEL - DOCIVYX</u>						
N 215813	002	10398785	Mar 14, 2036	DP		
<u>DOCETAXEL - DOCIVYX</u>						
N 215813	003	10398785	Mar 14, 2036	DP		
<u>DOCETAXEL - BEIZRAY</u>						
N 218711	001	11419842	May 16, 2036	DP U-4027		
		11419842	May 16, 2036	DP U-4028		
		11419842	May 16, 2036	DP U-881		
		11419842	May 16, 2036	DP U-937		
		11419842	May 16, 2036	DP U-946		
		12090134	May 16, 2036	DP U-4027		
		12090134	May 16, 2036	DP U-4028		
		12090134	May 16, 2036	DP U-881		
		12090134	May 16, 2036	DP U-937		
		12090134	May 16, 2036	DP U-946		
		12090135	May 16, 2036	DP		
<u>DOLUTEGRAVIR SODIUM - TIVICAY</u>						
N 204790	001	8129385	Oct 05, 2027	DS DP		
		8129385*PED	Apr 05, 2028			
		9242986	Dec 08, 2029	DS DP		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>DOLUTEGRAVIR SODIUM - TIVICAY</u>						
N 204790 001	9242986*PED	Jun 08, 2030				
<u>DOLUTEGRAVIR SODIUM - TIVICAY</u>						
N 204790 002	8129385	Oct 05, 2027	DS DP			
	8129385*PED	Apr 05, 2028				
	9242986	Dec 08, 2029	DS DP			
	9242986*PED	Jun 08, 2030				
<u>DOLUTEGRAVIR SODIUM - TIVICAY</u>						
N 204790 003	8129385	Oct 05, 2027	DS DP			
	8129385*PED	Apr 05, 2028				
	9242986	Dec 08, 2029	DS DP			
	9242986*PED	Jun 08, 2030				
<u>DOLUTEGRAVIR SODIUM - TIVICAY PD</u>						
N 213983 001	8129385	Oct 05, 2027	DS DP			
	8129385*PED	Apr 05, 2028				
	9242986	Dec 08, 2029	DS DP			
	9242986*PED	Jun 08, 2030				
<u>DOLUTEGRAVIR SODIUM; LAMIVUDINE - DOVATO</u>						
N 211994 001	11234985	Jan 24, 2031	DP U-257		NPP	Apr 05, 2027
	8129385	Oct 05, 2027	DS DP			
	8129385*PED	Apr 05, 2028				
	9242986	Dec 08, 2029	DS DP			
	9242986*PED	Jun 08, 2030				
<u>DOLUTEGRAVIR SODIUM; RILPIVIRINE HYDROCHLORIDE - JULUCA</u>						
N 210192 001	10426780	Jan 24, 2031	DS DP U-257			
	12011506	Sep 05, 2038	DP			
	7125879	Apr 21, 2025	DS DP U-257			
	8129385	Oct 05, 2027	DS DP			
	8129385*PED	Apr 05, 2028				
	9242986	Dec 08, 2029	DS DP			
	9242986*PED	Jun 08, 2030				
<u>DONEPEZIL HYDROCHLORIDE - ARICEPT</u>						
N 022568 001	8481565	Oct 04, 2026	DP			
<u>DONEPEZIL HYDROCHLORIDE - ADLARITY</u>						
N 212304 001	10016372	Jul 26, 2037	U-3334		NP	Mar 11, 2025
	10300025	Jul 26, 2037	DP			
	10307379	Jul 26, 2037	DP			
	10835499	May 20, 2038	DP U-3334			
	10966936	Aug 11, 2038	DP U-3334			
	11103463	Jul 26, 2037	U-3334			
	11648214	Sep 23, 2037	DP U-3334			
	11679086	May 26, 2037	U-3334			
	9993466	Jul 26, 2037	DP			
<u>DONEPEZIL HYDROCHLORIDE - ADLARITY</u>						
N 212304 002	10016372	Jul 26, 2037	U-3334		NP	Mar 11, 2025
	10300025	Jul 26, 2037	DP			
	10307379	Jul 26, 2037	DP			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>DONEPEZIL HYDROCHLORIDE - ADLARITY</u>						
N 212304 002	10835499	May 20, 2038	DP U-3334			
	10966936	Aug 11, 2038	DP U-3334			
	11103463	Jul 26, 2037	U-3334			
	11648214	Sep 23, 2037	DP U-3334			
	11679086	May 26, 2037	U-3334			
	9993466	Jul 26, 2037	DP			
<u>DONEPEZIL HYDROCHLORIDE; MEMANTINE HYDROCHLORIDE - NAMZARIC</u>						
N 206439 001	8039009	Mar 24, 2029	U-1641			
	8039009*PED	Sep 24, 2029				
	8058291	Dec 05, 2029	U-1641			
	8168209	Nov 22, 2025	DP	Y		
	8168209*PED	May 22, 2026				
	8173708	Nov 22, 2025	U-1641	Y		
	8173708*PED	May 22, 2026				
	8283379	Nov 22, 2025	U-1641	Y		
	8283379*PED	May 22, 2026				
	8293794	Nov 22, 2025	DP			
	8338485	Nov 22, 2025	DP			
	8338486	Nov 22, 2025	U-1641			
	8580858	Nov 22, 2025	U-1641			
<u>DONEPEZIL HYDROCHLORIDE; MEMANTINE HYDROCHLORIDE - NAMZARIC</u>						
N 206439 002	8039009	Mar 24, 2029	U-1641			
	8039009*PED	Sep 24, 2029				
	8058291	Dec 05, 2029	U-1641			
	8168209	Nov 22, 2025	DP	Y		
	8168209*PED	May 22, 2026				
	8173708	Nov 22, 2025	U-1641	Y		
	8173708*PED	May 22, 2026				
	8283379	Nov 22, 2025	U-1641	Y		
	8283379*PED	May 22, 2026				
	8293794	Nov 22, 2025	DP			
	8329752	Nov 22, 2025	DP	Y		
	8329752*PED	May 22, 2026				
	8338485	Nov 22, 2025	DP			
	8338486	Nov 22, 2025	U-1641			
	8362085	Nov 22, 2025	U-1641	Y		
	8362085*PED	May 22, 2026				
	8580858	Nov 22, 2025	U-1641			
	8598233	Nov 22, 2025	DP	Y		
	8598233*PED	May 22, 2026				
<u>DONEPEZIL HYDROCHLORIDE; MEMANTINE HYDROCHLORIDE - NAMZARIC</u>						
N 206439 003	8039009	Mar 24, 2029	U-1641			
	8039009*PED	Sep 24, 2029				
	8058291	Dec 05, 2029	U-1641			
	8293794	Nov 22, 2025	DP			
	8338485	Nov 22, 2025	DP			
	8338486	Nov 22, 2025	U-1641			
	8580858	Nov 22, 2025	U-1641			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>DONEPEZIL HYDROCHLORIDE; MEMANTINE HYDROCHLORIDE - NAMZARIC</u>						
N 206439 004	8039009	Mar 24, 2029	U-1641			
	8039009*PED	Sep 24, 2029				
	8058291	Dec 05, 2029	U-1641			
	8168209	Nov 22, 2025	DP	Y		
	8168209*PED	May 22, 2026				
	8173708	Nov 22, 2025	U-1641	Y		
	8173708*PED	May 22, 2026				
	8283379	Nov 22, 2025	U-1641	Y		
	8283379*PED	May 22, 2026				
	8293794	Nov 22, 2025	DP			
	8329752	Nov 22, 2025	DP	Y		
	8329752*PED	May 22, 2026				
	8338485	Nov 22, 2025	DP			
	8338486	Nov 22, 2025	U-1641			
	8362085	Nov 22, 2025	U-1641	Y		
	8362085*PED	May 22, 2026				
	8580858	Nov 22, 2025	U-1641			
	8598233	Nov 22, 2025	DP	Y		
	8598233*PED	May 22, 2026				
<u>DORAVIRINE - PIFELTRO</u>						
N 210806 001	8486975	Aug 30, 2032	DS DP U-2394		NPP	Jan 27, 2025
	8486975	Aug 30, 2032	DS DP U-2630			
	8486975	Aug 30, 2032	DS DP U-3308			
<u>DORAVIRINE; LAMIVUDINE; TENOFOVIR DISOPROXIL FUMARATE - DELSTRIGO</u>						
N 210807 001	10603282	Nov 29, 2036	DP		NPP	Jan 27, 2025
	10842751	Nov 29, 2036	DP			
	8486975	Aug 30, 2032	DS DP U-2395			
	8486975	Aug 30, 2032	DS DP U-2629			
	8486975	Aug 30, 2032	DS DP U-3307			
<u>DOXEPIN HYDROCHLORIDE - SILENOR</u>						
N 022036 001	10238620	May 18, 2027	U-620			
	10548871	Apr 11, 2028	U-620			
	10653660	Jul 20, 2027	U-620			
	10653662	May 18, 2027	U-620			
	11096920	Apr 11, 2028	U-620			
	11110074	Jul 20, 2027	U-620			
	11234954	Jan 18, 2028	U-620			
	12083090	May 18, 2027	U-620			
	7915307	Aug 24, 2027	U-620			
	8513299	Sep 07, 2030	U-620			
	9107898	May 01, 2028	U-620			
	9486437	May 18, 2027	U-620			
	9532971	Jun 01, 2029	DP			
	9572814	Jul 20, 2027	U-620			
	9861607	May 18, 2027	U-620			
	9907780	Apr 11, 2028	DP			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>DOXEPIH HYDROCHLORIDE - SILENOR</u>						
N 022036	002 10238620	May 18, 2027	U-620			
	10548871	Apr 11, 2028	U-620			
	10653660	Jul 20, 2027	U-620			
	10653662	May 18, 2027	U-620			
	11096920	Apr 11, 2028	U-620			
	11110074	Jul 20, 2027	U-620			
	11234954	Jan 18, 2028	U-620			
	12083090	May 18, 2027	U-620			
	7915307	Aug 24, 2027	U-620			
	8513299	Sep 07, 2030	U-620			
	9107898	May 01, 2028	U-620			
	9486437	May 18, 2027	U-620			
	9532971	Jun 01, 2029	DP			
	9572814	Jul 20, 2027	U-620			
	9861607	May 18, 2027	U-620			
	9907780	Apr 11, 2028	DP			
<u>DOXYCYCLINE - ORACEA</u>						
N 050805	001 7749532	Dec 19, 2027	DP U-1063			
	8206740	Dec 24, 2025	DP U-925			
<u>DOXYCYCLINE HYCLATE - DORYX</u>						
N 050795	001 8715724	Feb 03, 2028	DP			
<u>DOXYCYCLINE HYCLATE - DORYX</u>						
N 050795	002 8715724	Feb 03, 2028	DP			
<u>DOXYCYCLINE HYCLATE - DORYX</u>						
N 050795	003 8715724	Feb 03, 2028	DP			
<u>DOXYCYCLINE HYCLATE - DORYX</u>						
N 050795	004 8715724	Feb 03, 2028	DP			
<u>DOXYCYCLINE HYCLATE - DORYX</u>						
N 050795	005 8715724	Feb 03, 2028	DP			
<u>DOXYCYCLINE HYCLATE - DORYX</u>						
N 050795	006 8715724	Feb 03, 2028	DP			
<u>DOXYCYCLINE HYCLATE - DORYX MPC</u>						
N 050795	007 8715724	Feb 03, 2028	DP			
	9295652	Oct 23, 2034	DP U-918			
	9446057	Dec 23, 2034	DP U-918			
	9511031	Oct 23, 2034	DP			
<u>DOXYCYCLINE HYCLATE - DORYX MPC</u>						
N 050795	008 8715724	Feb 03, 2028	DP			
	9295652	Oct 23, 2034	DP U-918			
	9446057	Dec 23, 2034	DP U-918			
	9511031	Oct 23, 2034	DP			
<u>DOXYLAMINE SUCCINATE; PYRIDOXINE HYDROCHLORIDE - BONJESTA</u>						
N 209661	001 9089489	Feb 18, 2033	DP U-1382			
	9375404	Feb 18, 2033	DP U-1382			
	9526703	Feb 18, 2033	DP U-1382			
	9937132	Feb 18, 2033	DP U-1382			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>DOXYLAMINE SUCCINATE; PYRIDOXINE HYDROCHLORIDE - BONJESTA</u>						
N 209661	001 9089489	Feb 18, 2033	DP U-1382			
	9375404	Feb 18, 2033	DP U-1382			
	9526703	Feb 18, 2033	DP U-1382			
	9937132	Feb 18, 2033	DP U-1382			
<u>DRONABINOL - SYNDROS</u>						
N 205525	001 10265293	Aug 06, 2028	DS DP			
	11253472	Aug 06, 2028	DS DP			
	8222292	Aug 06, 2028	DS DP			
	9345771	Aug 06, 2028	DS DP			
<u>DRONEDARONE HYDROCHLORIDE - MULTAQ</u>						
N 022425	001 8410167	Apr 16, 2029	U-1387			
	8410167	Apr 16, 2029	U-1388			
	8602215	Jun 30, 2031	U-1473			
	9107900	Apr 16, 2029	U-1726			
	9107900	Apr 16, 2029	U-1728			
<u>DROSPIRENONE - SLYND</u>						
N 211367	001 10179140	Jun 28, 2031	U-2553			
	10603281	Jun 28, 2031	U-2553			
	10849857	Jun 28, 2031	DP U-2553			
	10987364	Jun 28, 2031	DP			
	11123299	Jun 28, 2031	DP			
	11291632	Jun 28, 2031	DP			
	11291633	Jun 28, 2031	DP			
	11351122	Jun 28, 2031	DP			
	11413249	Jun 28, 2031	U-2553			
	11439598	Jun 28, 2031	DP			
	11478487	Jun 28, 2031	DP			
	11504334	Jun 28, 2031	DP			
	11951213	Jun 28, 2031	U-2553			
	12090231	Jun 28, 2031	U-2553			
	9603860	Jun 28, 2031	U-2553			
<u>DROSPIRENONE - DROSPIRENONE</u>						
N 216285	001 10179140	Jun 28, 2031	U-2553		NP	Jun 29, 2025
	10603281	Jun 28, 2031	U-2553			
	10849857	Jun 28, 2031	DP U-2553			
	10987364	Jun 28, 2031	DP			
	11123299	Jun 28, 2031	DP			
	11291632	Jun 28, 2031	DP			
	11291633	Jun 28, 2031	DP			
	11351122	Jun 28, 2031	DP			
	11413249	Jun 28, 2031	U-2553			
	11439598	Jun 28, 2031	DP			
	11478487	Jun 28, 2031	DP			
	11504334	Jun 28, 2031	DP			
	11951213	Jun 28, 2031	U-2553			
	12090231	Jun 28, 2031	U-2553			
	9603860	Jun 28, 2031	U-2553			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>DROSPIRENONE; ESTETROL - NEXTSTELLIS</u>						
N 214154	001	11793760	Jun 17, 2036	DP	NCE	Apr 15, 2026
		11957694	Jun 17, 2036	DP		
		11964055	Jun 17, 2036	DP		
		7732430	Mar 02, 2026	DP U-3152		
<u>DROSPIRENONE; ESTRADIOL - ANGELIO</u>						
N 021355	001	8906890	Oct 22, 2031	DP		
<u>DROSPIRENONE; ETHINYL ESTRADIOL; LEVOMEFOLATE CALCIUM - BEYAZ</u>						
N 022532	001	11617751	Jul 17, 2030	DP U-1		
		11617751	Jul 17, 2030	DP U-3572		
		11617751	Jul 17, 2030	DP U-3573		
		11617751	Jul 17, 2030	DP U-3574		
		8617597	Feb 08, 2030	DP		
<u>DROSPIRENONE; ETHINYL ESTRADIOL; LEVOMEFOLATE CALCIUM - SAFYRAL</u>						
N 022574	001	11617751	Jul 17, 2030	DP U-1		
		11617751	Jul 17, 2030	DP U-3572		
		8617597	Feb 08, 2030	DP		
<u>DULOXETINE HYDROCHLORIDE - DRIZALMA SPRINKLE</u>						
N 212516	001	10413525	Apr 13, 2037	DP		
		10959982	Apr 13, 2037	DP		
		11202772	Apr 13, 2037	DP		
		9839626	Apr 13, 2037	DP		
<u>DULOXETINE HYDROCHLORIDE - DRIZALMA SPRINKLE</u>						
N 212516	002	10413525	Apr 13, 2037	DP		
		10959982	Apr 13, 2037	DP		
		11202772	Apr 13, 2037	DP		
		9839626	Apr 13, 2037	DP		
<u>DULOXETINE HYDROCHLORIDE - DRIZALMA SPRINKLE</u>						
N 212516	003	10413525	Apr 13, 2037	DP		
		10959982	Apr 13, 2037	DP		
		11202772	Apr 13, 2037	DP		
		9839626	Apr 13, 2037	DP		
<u>DULOXETINE HYDROCHLORIDE - DRIZALMA SPRINKLE</u>						
N 212516	004	10413525	Apr 13, 2037	DP		
		10959982	Apr 13, 2037	DP		
		11202772	Apr 13, 2037	DP		
		9839626	Apr 13, 2037	DP		
<u>DURLOBACTAM SODIUM; DURLOBACTAM SODIUM; SULBACTAM SODIUM - XACDURO (COPACKAGED)</u>						
N 216974	001	10376499	Nov 17, 2035	DP U-2840	NCE	May 23, 2028
		9309245	Apr 02, 2033	DS	GAIN	May 23, 2033
		9623014	Apr 02, 2033	U-2840		
		9968593	Nov 17, 2035	DP U-2840		
<u>DUVELISIB - COPIKTRA</u>						
N 211155	001	11312718	Jan 10, 2032	DP	ODE-208	Sep 24, 2025
		8193182	Feb 13, 2030	DS		
		9216982	Jan 05, 2029	U-2412		
		9216982	Jan 05, 2029	U-2413		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>DUVELISIB - COPIKTRA</u>						
N 211155	001	9840505	Jan 10, 2032	U-2412		
		9840505	Jan 10, 2032	U-2413		
		RE46621	May 17, 2032	DS DP		
<u>DUVELISIB - COPIKTRA</u>						
N 211155	002	11312718	Jan 10, 2032	DP	ODE-208	Sep 24, 2025
		8193182	Feb 13, 2030	DS		
		9216982	Jan 05, 2029	U-2412		
		9216982	Jan 05, 2029	U-2413		
		9840505	Jan 10, 2032	U-2412		
		9840505	Jan 10, 2032	U-2413		
		RE46621	May 17, 2032	DS DP		
<u>ECONAZOLE NITRATE - ECOZA</u>						
N 205175	001	10071054	Aug 08, 2031	DP		
<u>EDARAVONE - RADICAVA ORS</u>						
N 215446	001	10987341	Nov 01, 2039	DP	NP	May 12, 2025
		11241416	Nov 01, 2039	DP	ODE-144	May 12, 2029
		11478450	Nov 01, 2039	U-3468		
		11826352	Nov 01, 2039	DP		
		11957660	Nov 01, 2039	DP		
<u>EDOXABAN TOSYLATE - SAVAYSA</u>						
N 206316	001	7365205	Apr 18, 2027	DS	M-14	Oct 18, 2026
		9149532	Mar 28, 2028	DP		
<u>EDOXABAN TOSYLATE - SAVAYSA</u>						
N 206316	002	7365205	Apr 18, 2027	DS	M-14	Oct 18, 2026
		9149532	Mar 28, 2028	DP		
<u>EDOXABAN TOSYLATE - SAVAYSA</u>						
N 206316	003	7365205	Apr 18, 2027	DS	M-14	Oct 18, 2026
		9149532	Mar 28, 2028	DP		
<u>EFAVIRENZ; EMTRICITABINE; TENOFOVIR DISOPROXIL FUMARATE - ATRIPLA</u>						
N 021937	001	8598185	Apr 28, 2029	DP		
		9018192	Jun 13, 2026	U-1170		
		9018192	Jun 13, 2026	U-750		
		9545414	Jun 13, 2026	DP U-1170		
		9545414	Jun 13, 2026	DP U-750		
<u>EFINACONAZOLE - JUBLIA</u>						
N 203567	001	10105444	Jul 08, 2030	DP		
		10342875	Oct 02, 2034	DP U-2720		
		10478601	Apr 25, 2035	DP U-2721		
		10512640	Jan 03, 2028	U-1969		
		10828293	Oct 02, 2034	U-2720		
		10828369	Jan 03, 2028	DP		
		10864274	Oct 02, 2034	U-2720		
		11213519	Jan 03, 2028	U-2720		
		11654139	Oct 02, 2034	U-1969		
		11872218	Jan 03, 2028	U-1969		
		7214506	Feb 22, 2026	U-281		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>EFINACONAZOLE - JUBLIA</u>						
N 203567	001 8039494	Jul 08, 2030	U-281			
	8486978	Oct 24, 2030	DP			
	9302009	Oct 24, 2030	DP			
	9566272	Jan 03, 2028	U-1969			
	9662394	Oct 02, 2034	DP			
	9861698	Jul 08, 2030	DP			
	9877955	Jan 03, 2028	U-1969			
<u>EFLORNITHINE HYDROCHLORIDE - IWILFIN</u>						
N 215500	001				NP	Dec 13, 2026
					ODE-462	Dec 13, 2030
<u>ELACESTRANT HYDROCHLORIDE - ORSERDU</u>						
N 217639	001 10071066	Oct 10, 2034	U-3524		NCE	Jan 27, 2028
	10385008	Jan 05, 2038	DS DP			
	10420734	Oct 10, 2034	U-3524			
	10745343	Jan 05, 2038	U-3523			
	11779552	Oct 10, 2034	U-3524			
	11819480	Nov 29, 2036	U-3523			
	7612114	Aug 18, 2026	DS DP U-3523			
	8399520	Dec 25, 2025	DS DP U-3523			
<u>ELACESTRANT HYDROCHLORIDE - ORSERDU</u>						
N 217639	002 10071066	Oct 10, 2034	U-3524		NCE	Jan 27, 2028
	10385008	Jan 05, 2038	DS DP			
	10420734	Oct 10, 2034	U-3524			
	10745343	Jan 05, 2038	U-3523			
	11779552	Oct 10, 2034	U-3524			
	11819480	Nov 29, 2036	U-3523			
	7612114	Aug 18, 2026	DS DP U-3523			
	8399520	Dec 25, 2025	DS DP U-3523			
<u>ELAFIBRANOR - IQIRVO</u>						
N 218860	001 11185519	Mar 30, 2037	U-3955		NCE	Jun 10, 2029
	11331292	Mar 30, 2037	U-1854		ODE-486	Jun 10, 2031
	11850223	Mar 30, 2037	U-1854			
	11857523	Mar 30, 2037	U-3955			
<u>ELAGOLIX SODIUM - ORILISSA</u>						
N 210450	001 10537572	Sep 01, 2036	U-2735			
	10682351	Sep 01, 2036	U-2850			
	11542239	Jul 23, 2039	DS DP			
	11690845	Aug 27, 2040	U-3654			
	11690854	Apr 19, 2038	U-3653			
	11707464	Mar 14, 2034	U-3672			
	12102637	Aug 20, 2038	DP			
	7419983	Jul 06, 2029	DS DP U-2360			
<u>ELAGOLIX SODIUM - ORILISSA</u>						
N 210450	002 11344551	Mar 14, 2034	U-3388			
	11344551	Mar 14, 2034	U-3389			
	11542239	Jul 23, 2039	DS DP			
	11690845	Aug 27, 2040	U-3654			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>ELAGOLIX SODIUM - ORILISSA</u>						
N 210450	002 12102637	Aug 20, 2038	DP			
	7419983	Jul 06, 2029	DS DP U-2360			
<u>ELAGOLIX SODIUM, ESTRADIOL, NORETHINDRONE ACETATE; ELAGOLIX SODIUM - ORIAHNN (COPACKAGED)</u>						
N 213388	001 10881659	Mar 14, 2034	U-2842			
	11045470	Mar 14, 2034	U-2842			
	11459305	Nov 07, 2028	DP U-2842			
	11542239	Jul 23, 2039	DS DP			
	11690845	Aug 27, 2040	U-3655			
	12083227	Aug 20, 2038	DP U-2842			
	7419983	Jul 06, 2029	DS DP			
<u>ELBASVIR; GRAZOPREVRIL - ZEPATIER</u>						
N 208261	001 7973040	Jul 24, 2029	DS DP U-1813			
	8871759	May 04, 2031	DS DP U-1813			
<u>ELEXACAFTOR, IVACAFTOR, TEZACAFTOR; IVACAFTOR - TRIKAFTA (COPACKAGED)</u>						
N 212273	001 10022352	Apr 09, 2027	DP U-2651		ODE-275	Oct 21, 2026
	10022352	Apr 09, 2027	DP U-3156		ODE-323	Dec 21, 2027
	10081621	Mar 25, 2031	DP U-2652		ODE-357	Jun 08, 2028
	10081621	Mar 25, 2031	DP U-3032			
	10081621	Mar 25, 2031	DP U-3157			
	10239867	Apr 09, 2027	DS DP U-2653			
	10239867	Apr 09, 2027	DS DP U-3033			
	10239867	Apr 09, 2027	DS DP U-3158			
	10646481	Aug 13, 2029	DP			
	10758534	Oct 06, 2035	DS DP U-2645			
	10758534	Oct 06, 2035	DS DP U-3028			
	10758534	Oct 06, 2035	DS DP U-3144			
	10793547	Dec 08, 2037	DS DP U-2645			
	10793547	Dec 08, 2037	DS DP U-3028			
	10793547	Dec 08, 2037	DS DP U-3144			
	11179367	Dec 08, 2037	DP U-3253			
	11426407	Oct 06, 2035	DS DP U-3425			
	11453655	Dec 08, 2037	DS DP			
	11517564	Dec 08, 2037	DP U-3498			
	11564916	Aug 13, 2029	U-3525			
	11578062	Mar 25, 2031	DP U-3544			
	11639347	Apr 09, 2027	DS DP U-3587			
	7495103	May 20, 2027	DS DP			
	7645789	May 01, 2027	DS DP			
	7776905	Jun 03, 2027	DS DP			
	8324242	Aug 05, 2027	U-2645			
	8324242	Aug 05, 2027	U-3028			
	8324242	Aug 05, 2027	U-3144			
	8354427	Jul 06, 2026	U-3029			
	8354427	Jul 06, 2026	U-3145			
	8410274	Dec 28, 2026	DP			
	8415387	Nov 12, 2027	U-2645			
	8415387	Nov 12, 2027	U-3028			
	8415387	Nov 12, 2027	U-3144			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>ELEXACAFITOR, IVACAFITOR, TEZACAFITOR; IVACAFITOR - TRIKAFTA (COPACKAGED)</u>						
N 212273 001	8598181	May 01, 2027	U-2645			
	8598181	May 01, 2027	U-3028			
	8598181	May 01, 2027	U-3144			
	8623905	May 01, 2027	DS DP			
	8629162	Jun 24, 2025	U-2648			
	8629162	Jun 24, 2025	U-3030			
	8629162	Jun 24, 2025	U-3146			
	8754224	Dec 28, 2026	DS DP			
	9012496	Jul 15, 2033	U-2649			
	9012496	Jul 15, 2033	U-3154			
	9670163	Dec 28, 2026	DP U-2650			
	9670163	Dec 28, 2026	DP U-3031			
	9670163	Dec 28, 2026	DP U-3155			
	9931334	Dec 28, 2026	DP U-2650			
	9931334	Dec 28, 2026	DP U-3031			
	9931334	Dec 28, 2026	DP U-3155			
	9974781	Apr 09, 2027	DP U-2645			
	9974781	Apr 09, 2027	DP U-3028			
	9974781	Apr 09, 2027	DP U-3144			
<u>ELEXACAFITOR, IVACAFITOR, TEZACAFITOR; IVACAFITOR - TRIKAFTA (COPACKAGED)</u>						
N 212273 002	10022352	Apr 09, 2027	DP U-3156		ODE-357	Jun 08, 2028
	10081621	Mar 25, 2031	DP U-3157			
	10239867	Apr 09, 2027	DS DP U-3158			
	10646481	Aug 13, 2029	DP			
	10758534	Oct 06, 2035	DS DP U-3144			
	10793547	Dec 08, 2037	DS DP U-3144			
	11179367	Dec 08, 2037	DP U-3253			
	11426407	Oct 06, 2035	DS DP U-3425			
	11453655	Dec 08, 2037	DS DP			
	11517564	Dec 08, 2037	DP U-3498			
	11564916	Aug 13, 2029	U-3525			
	11578062	Mar 25, 2031	DP U-3544			
	11639347	Apr 09, 2027	DS DP U-3587			
	7495103	May 20, 2027	DS DP			
	7645789	May 01, 2027	DS DP			
	7776905	Jun 03, 2027	DS DP			
	8324242	Aug 05, 2027	U-3144			
	8354427	Jul 06, 2026	U-3145			
	8410274	Dec 28, 2026	DP			
	8415387	Nov 12, 2027	U-3144			
	8598181	May 01, 2027	U-3144			
	8623905	May 01, 2027	DS DP			
	8629162	Jun 24, 2025	U-3146			
	8754224	Dec 28, 2026	DS DP			
	9012496	Jul 15, 2033	U-3154			
	9670163	Dec 28, 2026	DP U-3155			
	9931334	Dec 28, 2026	DP U-3155			
	9974781	Apr 09, 2027	DP U-3144			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>ELEXACAFTOR, IVACAFTOR, TEZACAFTOR; IVACAFTOR - TRIKAFTA (COPACKAGED)</u>						
N 217660 001	10022352	Apr 09, 2027	DP U-3596		NP	Apr 26, 2026
	10081621	Mar 25, 2031	DP U-3600		ODE-433	Apr 26, 2030
	10239867	Apr 09, 2027	DS DP U-3590			
	10272046	Feb 27, 2033	DP U-3599			
	10646481	Aug 13, 2029	DP			
	10758534	Oct 06, 2035	DS DP U-3589			
	10793547	Dec 08, 2037	DS DP U-3588			
	11147770	Feb 27, 2033	DP U-3598			
	11179367	Dec 08, 2037	DP U-3597			
	11426407	Oct 06, 2035	DS DP U-3595			
	11453655	Dec 08, 2037	DS DP			
	11517564	Dec 08, 2037	DP U-3586			
	11564916	Aug 13, 2029	U-3585			
	11578062	Mar 25, 2031	DP U-3584			
	11639347	Apr 09, 2027	DS DP U-3583			
	11752106	Feb 27, 2033	DP U-3696			
	7495103	May 20, 2027	DS DP			
	7645789	May 01, 2027	DS DP			
	7776905	Jun 03, 2027	DS DP			
	8324242	Aug 05, 2027	U-3589			
	8354427	Jul 06, 2026	U-3593			
	8410274	Dec 28, 2026	DP			
	8415387	Nov 12, 2027	U-3589			
	8598181	May 01, 2027	U-3589			
	8623905	May 01, 2027	DS DP			
	8629162	Jun 24, 2025	U-3592			
	8754224	Dec 28, 2026	DS DP			
	8883206	Feb 27, 2033	DP			
	9670163	Dec 28, 2026	DP U-3591			
	9931334	Dec 28, 2026	DP U-3591			
	9974781	Apr 09, 2027	DP U-3589			
<u>ELEXACAFTOR, IVACAFTOR, TEZACAFTOR; IVACAFTOR - TRIKAFTA (COPACKAGED)</u>						
N 217660 002	10022352	Apr 09, 2027	DP U-3596		NP	Apr 26, 2026
	10081621	Mar 25, 2031	DP U-3600		ODE-433	Apr 26, 2030
	10239867	Apr 09, 2027	DS DP U-3590			
	10272046	Feb 27, 2033	DP U-3599			
	10646481	Aug 13, 2029	DP			
	10758534	Oct 06, 2035	DS DP U-3589			
	10793547	Dec 08, 2037	DS DP U-3588			
	11147770	Feb 27, 2033	DP U-3598			
	11179367	Dec 08, 2037	DP U-3597			
	11426407	Oct 06, 2035	DS DP U-3595			
	11453655	Dec 08, 2037	DS DP			
	11517564	Dec 08, 2037	DP U-3586			
	11564916	Aug 13, 2029	U-3585			
	11578062	Mar 25, 2031	DP U-3584			
	11639347	Apr 09, 2027	DS DP U-3583			
	11752106	Feb 27, 2033	DP U-3696			
	7495103	May 20, 2027	DS DP			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>ELEXACAFTOR, IVACAFTOR, TEZACAFTOR; IVACAFTOR - TRIKAFTA (COPACKAGED)</u>						
N 217660 002	7645789	May 01, 2027	DS DP			
	7776905	Jun 03, 2027	DS DP			
	8324242	Aug 05, 2027		U-3589		
	8354427	Jul 06, 2026		U-3593		
	8410274	Dec 28, 2026	DP			
	8415387	Nov 12, 2027		U-3589		
	8598181	May 01, 2027		U-3589		
	8623905	May 01, 2027	DS DP			
	8629162	Jun 24, 2025		U-3592		
	8754224	Dec 28, 2026	DS DP			
	8883206	Feb 27, 2033	DP			
	9670163	Dec 28, 2026	DP	U-3591		
	9931334	Dec 28, 2026	DP	U-3591		
	9974781	Apr 09, 2027	DP	U-3589		
<u>ELIPLUSTAT TARTRATE - CERDELGA</u>						
N 205494 001	10888544	Dec 13, 2038		U-3040		
	10888544	Dec 13, 2038		U-3041		
	10888547	Jan 31, 2031		U-3042		
	10888547	Jan 31, 2031		U-3043		
	11458119	Nov 24, 2030	DS DP			
	7196205	Jun 26, 2026	DS			
<u>ELTROMBOPAG CHOLINE - ALVAIZ</u>						
N 216774 001	11072586	Nov 05, 2038	DS DP	U-3936		
	11072586	Nov 05, 2038	DS DP	U-3937		
	11072586	Nov 05, 2038	DS DP	U-3938		
<u>ELTROMBOPAG CHOLINE - ALVAIZ</u>						
N 216774 002	11072586	Nov 05, 2038	DS DP	U-3936		
	11072586	Nov 05, 2038	DS DP	U-3937		
	11072586	Nov 05, 2038	DS DP	U-3938		
<u>ELTROMBOPAG CHOLINE - ALVAIZ</u>						
N 216774 003	11072586	Nov 05, 2038	DS DP	U-3936		
	11072586	Nov 05, 2038	DS DP	U-3937		
	11072586	Nov 05, 2038	DS DP	U-3938		
<u>ELTROMBOPAG CHOLINE - ALVAIZ</u>						
N 216774 004	11072586	Nov 05, 2038	DS DP	U-3936		
	11072586	Nov 05, 2038	DS DP	U-3937		
	11072586	Nov 05, 2038	DS DP	U-3938		
<u>ELTROMBOPAG OLAMINE - PROMACTA</u>						
N 022291 001	7547719	Jul 13, 2025	DS DP	U-1306	ODE-210	Nov 16, 2025
	7547719	Jul 13, 2025	DS DP	U-1575		
	7547719	Jul 13, 2025	DS DP	U-2451		
	7547719	Jul 13, 2025	DS DP	U-2452		
	7547719*PED	Jan 13, 2026				
	8052993	Aug 01, 2027	DP	U-1306		
	8052993	Aug 01, 2027	DP	U-1575		
	8052993	Aug 01, 2027	DP	U-2451		
	8052993	Aug 01, 2027	DP	U-930		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>ELTROMBOPAG OLAMINE - PROMACTA</u>						
N 022291 001	8052993*PED	Feb 01, 2028				
	8052994	Aug 01, 2027	DP U-1714			
	8052994*PED	Feb 01, 2028				
	8062665	Aug 01, 2027	DP U-1714			
	8062665*PED	Feb 01, 2028				
	8071129	Aug 01, 2027	DP U-1714			
	8071129*PED	Feb 01, 2028				
	8828430	Aug 01, 2027	DP U-1306			
	8828430	Aug 01, 2027	DP U-2451			
	8828430*PED	Feb 01, 2028				
<u>ELTROMBOPAG OLAMINE - PROMACTA</u>						
N 022291 002	7547719	Jul 13, 2025	DS DP U-1306		ODE-210	Nov 16, 2025
	7547719	Jul 13, 2025	DS DP U-1575			
	7547719	Jul 13, 2025	DS DP U-2451			
	7547719	Jul 13, 2025	DS DP U-2452			
	7547719	Jul 13, 2025	DS DP U-930			
	7547719*PED	Jan 13, 2026				
	8052993	Aug 01, 2027	DP U-1714			
	8052993*PED	Feb 01, 2028				
	8052994	Aug 01, 2027	DP U-1306			
	8052994	Aug 01, 2027	DP U-2451			
	8052994	Aug 01, 2027	DP U-930			
	8052994*PED	Feb 01, 2028				
	8062665	Aug 01, 2027	DP U-1714			
	8062665*PED	Feb 01, 2028				
	8071129	Aug 01, 2027	DP U-1714			
	8071129*PED	Feb 01, 2028				
	8828430	Aug 01, 2027	DP U-1306			
	8828430	Aug 01, 2027	DP U-2451			
	8828430*PED	Feb 01, 2028				
<u>ELTROMBOPAG OLAMINE - PROMACTA</u>						
N 022291 003	7547719	Jul 13, 2025	DS DP U-1306		ODE-210	Nov 16, 2025
	7547719	Jul 13, 2025	DS DP U-1575			
	7547719	Jul 13, 2025	DS DP U-2451			
	7547719	Jul 13, 2025	DS DP U-2452			
	7547719	Jul 13, 2025	DS DP U-930			
	7547719*PED	Jan 13, 2026				
	8052993	Aug 01, 2027	DP U-1714			
	8052993*PED	Feb 01, 2028				
	8052994	Aug 01, 2027	DP U-1714			
	8052994*PED	Feb 01, 2028				
	8062665	Aug 01, 2027	DP U-1306			
	8062665	Aug 01, 2027	DP U-2451			
	8062665	Aug 01, 2027	DP U-930			
	8062665*PED	Feb 01, 2028				
	8071129	Aug 01, 2027	DP U-1714			
	8071129*PED	Feb 01, 2028				
	8828430	Aug 01, 2027	DP U-1306			
	8828430	Aug 01, 2027	DP U-2451			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>ELTROMBOPAG OLAMINE - PROMACTA</u>						
N 022291 003	8828430*PED	Feb 01, 2028				
<u>ELTROMBOPAG OLAMINE - PROMACTA</u>						
N 022291 004	7547719	Jul 13, 2025	DS DP U-1306		ODE-210	Nov 16, 2025
	7547719	Jul 13, 2025	DS DP U-1575			
	7547719	Jul 13, 2025	DS DP U-2451			
	7547719	Jul 13, 2025	DS DP U-2452			
	7547719*PED	Jan 13, 2026				
	8052993	Aug 01, 2027	DP U-1714			
	8052993*PED	Feb 01, 2028				
	8052994	Aug 01, 2027	DP U-1714			
	8052994*PED	Feb 01, 2028				
	8062665	Aug 01, 2027	DP U-1714			
	8062665*PED	Feb 01, 2028				
	8071129	Aug 01, 2027	DP U-1306			
	8071129	Aug 01, 2027	DP U-2451			
	8071129	Aug 01, 2027	DP U-930			
	8071129*PED	Feb 01, 2028				
	8828430	Aug 01, 2027	DP U-1306			
	8828430	Aug 01, 2027	DP U-2451			
	8828430*PED	Feb 01, 2028				
<u>ELTROMBOPAG OLAMINE - PROMACTA</u>						
N 022291 005	7547719	Jul 13, 2025	DS DP U-1306		ODE-210	Nov 16, 2025
	7547719	Jul 13, 2025	DS DP U-1575			
	7547719	Jul 13, 2025	DS DP U-930			
	7547719*PED	Jan 13, 2026				
<u>ELTROMBOPAG OLAMINE - PROMACTA KIT</u>						
N 207027 001	7547719	Jul 13, 2025	DS DP U-1306		ODE*	Nov 16, 2025
	7547719	Jul 13, 2025	DS DP U-1575			
	7547719	Jul 13, 2025	DS DP U-1736			
	7547719	Jul 13, 2025	DS DP U-2452			
	7547719*PED	Jan 13, 2026				
<u>ELTROMBOPAG OLAMINE - PROMACTA KIT</u>						
N 207027 002	7547719	Jul 13, 2025	DS DP U-1306		ODE*	Nov 16, 2025
	7547719	Jul 13, 2025	DS DP U-1575			
	7547719	Jul 13, 2025	DS DP U-1736			
	7547719	Jul 13, 2025	DS DP U-2452			
	7547719*PED	Jan 13, 2026				
<u>ELUXADOLINE - VIBERZI</u>						
N 206940 001	10188632	Mar 14, 2033	DP			
	10213415	Mar 14, 2025	DS U-2152			
	11007179	Mar 14, 2033	DP			
	11090291	Mar 14, 2033	DP			
	11160792	Mar 14, 2033	DP			
	11229627	Mar 14, 2033	DP			
	11311516	Mar 14, 2033	DP			
	12097187	Mar 14, 2033	U-3475			
	7741356	May 27, 2029	DS DP			
	7786158	Mar 14, 2025	DS			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>ELUXADOLINE - VIBERZI</u>						
N 206940 001	8344011	Mar 14, 2025	U-1709			
	8609709	Mar 14, 2025	DS			
	8691860	Jul 07, 2028	DS	U-1709		
	8772325	Mar 14, 2025	U-1709			
	9115091	Jul 07, 2028	DS DP	U-1738		
	9205076	Mar 14, 2025	U-1709			
	9364489	Jul 07, 2028	U-1709			
	9675587	Mar 14, 2033	DP			
	9700542	Mar 14, 2025	DP			
	9789125	Jul 07, 2028	DP	U-1709		
	9789125	Jul 07, 2028	DP	U-2152		
<u>ELUXADOLINE - VIBERZI</u>						
N 206940 002	10188632	Mar 14, 2033	DP			
	10213415	Mar 14, 2025	DS	U-2152		
	11007179	Mar 14, 2033	DP			
	11090291	Mar 14, 2033	DP			
	11160792	Mar 14, 2033	DP			
	11229627	Mar 14, 2033	DP			
	11311516	Mar 14, 2033	DP			
	11484527	Mar 14, 2033	U-3475			
	7741356	May 27, 2029	DS DP			
	7786158	Mar 14, 2025	DS			
	8344011	Mar 14, 2025	U-1709			
	8609709	Mar 14, 2025	DS			
	8691860	Jul 07, 2028	DS	U-1709		
	8772325	Mar 14, 2025	U-1709			
	9115091	Jul 07, 2028	DS DP	U-1738		
	9205076	Mar 14, 2025	U-1709			
	9364489	Jul 07, 2028	U-1709			
	9675587	Mar 14, 2033	DP			
	9700542	Mar 14, 2025	DP			
	9789125	Jul 07, 2028	DP	U-1709		
	9789125	Jul 07, 2028	DP	U-2152		
<u>ELVITEGRAVIR - VITEKTA</u>						
N 203093 001	7176220	Aug 27, 2026	DS DP	U-257		
	7176220*PED	Feb 27, 2027				
	7635704	Oct 26, 2026	DS DP	U-257		
	7635704*PED	Apr 26, 2027				
	8981103	Oct 26, 2026	DS DP			
	8981103*PED	Apr 26, 2027				
<u>ELVITEGRAVIR - VITEKTA</u>						
N 203093 002	7176220	Aug 27, 2026	DS DP	U-257		
	7176220*PED	Feb 27, 2027				
	7635704	Oct 26, 2026	DS DP	U-257		
	7635704*PED	Apr 26, 2027				
	8981103	Oct 26, 2026	DS DP			
	8981103*PED	Apr 26, 2027				

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>EMPAGLIFLOZIN - JARDIANCE</u>						
N 204629 001	10258637	Apr 03, 2034	U-2290		I-922	Sep 21, 2026
	10258637*PED	Oct 03, 2034			M-82	Feb 24, 2025
	11090323	Apr 03, 2034	U-3191		NPP	Jun 20, 2026
	11090323*PED	Oct 03, 2034			PED	Aug 24, 2025
	11666590	Apr 03, 2034	U-3691		PED	Dec 20, 2026
	11813275	Apr 03, 2034	U-3759			
	11813275	Apr 03, 2034	U-3760			
	11833166	Apr 03, 2034	U-3776			
	11833166	Apr 03, 2034	U-3777			
	12115179	Feb 11, 2030	U-4023			
	7579449	Aug 01, 2028	DS			
	7579449*PED	Feb 01, 2029				
	7713938	Apr 15, 2027	DS DP			
	7713938*PED	Oct 15, 2027				
	8551957	Oct 14, 2029	U-1651			
	8551957*PED	Apr 14, 2030				
	9949997	May 17, 2034	U-2292			
	9949997	May 17, 2034	U-3199			
	9949997	May 17, 2034	U-3325			
	9949997*PED	Nov 17, 2034				
	9949998	Jun 11, 2034	U-2290			
	9949998*PED	Dec 11, 2034				
<u>EMPAGLIFLOZIN - JARDIANCE</u>						
N 204629 002	10258637	Apr 03, 2034	U-2290		NPP	Jun 20, 2026
	10258637*PED	Oct 03, 2034			PED	Dec 20, 2026
	11090323	Apr 03, 2034	U-3191			
	11090323*PED	Oct 03, 2034				
	11833166	Apr 03, 2034	U-3776			
	11833166	Apr 03, 2034	U-3777			
	12115179	Feb 11, 2030	U-4023			
	7579449	Aug 01, 2028	DS			
	7579449*PED	Feb 01, 2029				
	7713938	Apr 15, 2027	DS DP			
	7713938*PED	Oct 15, 2027				
	8551957	Oct 14, 2029	U-1651			
	8551957*PED	Apr 14, 2030				
	9949997	May 17, 2034	U-2292			
	9949997*PED	Nov 17, 2034				
	9949998	Jun 11, 2034	U-2290			
	9949998*PED	Dec 11, 2034				
<u>EMPAGLIFLOZIN; LINAGLIPTIN - GLYXAMBI</u>						
N 206073 001	10258637	Apr 03, 2034	U-2290			
	10258637*PED	Oct 03, 2034				
	11033552	May 04, 2027	DP			
	11033552*PED	Nov 04, 2027				
	11090323	Apr 03, 2034	U-3191			
	11090323*PED	Oct 03, 2034				
	11833166	Apr 03, 2034	U-3776			
	11833166	Apr 03, 2034	U-3777			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>EMPAGLIFLOZIN; LINAGLIPTIN - GLYXAMBI</u>						
N 206073 001	12115179	Feb 11, 2030	U-4023			
	7407955	May 02, 2025	DS DP			
	7407955*PED	Nov 02, 2025				
	7579449	Aug 01, 2028	DS			
	7579449*PED	Feb 01, 2029				
	7713938	Apr 15, 2027	DS DP			
	7713938*PED	Oct 15, 2027				
	8551957	Oct 14, 2029	DP U-1651			
	8551957*PED	Apr 14, 2030				
	8673927	May 04, 2027	U-1652	Y		
	8673927*PED	Nov 04, 2027				
	8883805	Nov 26, 2025	DP			
	8883805*PED	May 26, 2026				
	9173859	May 04, 2027	DP U-1772	Y		
	9949998	Jun 11, 2034	U-2290			
<u>EMPAGLIFLOZIN; LINAGLIPTIN - GLYXAMBI</u>						
N 206073 002	10258637	Apr 03, 2034	U-2290			
	10258637*PED	Oct 03, 2034				
	11033552	May 04, 2027	DP			
	11033552*PED	Nov 04, 2027				
	11090323	Apr 03, 2034	U-3191			
	11090323*PED	Oct 03, 2034				
	11833166	Apr 03, 2034	U-3776			
	11833166	Apr 03, 2034	U-3777			
	12115179	Feb 11, 2030	U-4023			
	7407955	May 02, 2025	DS DP			
	7407955*PED	Nov 02, 2025				
	7579449	Aug 01, 2028	DS			
	7579449*PED	Feb 01, 2029				
	7713938	Apr 15, 2027	DS DP			
	7713938*PED	Oct 15, 2027				
	8551957	Oct 14, 2029	DP U-1651			
	8551957*PED	Apr 14, 2030				
	8673927	May 04, 2027	U-1652	Y		
	8673927*PED	Nov 04, 2027				
	8883805	Nov 26, 2025	DP			
	8883805*PED	May 26, 2026				
	9173859	May 04, 2027	DP U-1772	Y		
	9949998	Jun 11, 2034	U-2290			
<u>EMPAGLIFLOZIN; LINAGLIPTIN; METFORMIN HYDROCHLORIDE - TRIJARDY XR</u>						
N 212614 001	10022379	Apr 02, 2029	DP U-2732			
	10258637	Apr 03, 2034	U-2731			
	10258637*PED	Oct 03, 2034				
	10406172	Jun 15, 2030	DP U-2733			
	10596120	Mar 07, 2032	DP U-2776			
	10596120	Mar 07, 2032	DP U-2790			
	11090323	Apr 03, 2034	U-3192			
	11090323*PED	Oct 03, 2034				
	11564886	Mar 07, 2032	DP U-3531			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>EMPAGLIFLOZIN; LINAGLIPTIN; METFORMIN HYDROCHLORIDE - TRIJARDY XR</u>						
N 212614 001	11833166	Apr 03, 2034	U-3776			
	11833166	Apr 03, 2034	U-3777			
	12115179	Feb 11, 2030	U-4023			
	7407955	May 02, 2025	DS DP			
	7407955*PED	Nov 02, 2025				
	7579449	Aug 01, 2028	DS			
	7579449*PED	Feb 01, 2029				
	7713938	Apr 15, 2027	DS DP			
	7713938*PED	Oct 15, 2027				
	8551957	Oct 14, 2029	DP U-2730			
	8551957*PED	Apr 14, 2030				
	8883805	Nov 26, 2025	DP			
	8883805*PED	May 26, 2026				
	9155705	May 21, 2030	DP			
	9415016	Apr 02, 2029	DP			
	9949998	Jun 11, 2034	U-2731			
	9949998*PED	Dec 11, 2034				
<u>EMPAGLIFLOZIN; LINAGLIPTIN; METFORMIN HYDROCHLORIDE - TRIJARDY XR</u>						
N 212614 002	10022379	Apr 02, 2029	DP U-2732			
	10258637	Apr 03, 2034	U-2731			
	10258637*PED	Oct 03, 2034				
	10406172	Jun 15, 2030	DP U-2733			
	10596120	Mar 07, 2032	DP U-2776			
	10596120	Mar 07, 2032	DP U-2790			
	11090323	Apr 03, 2034	U-3192			
	11090323*PED	Oct 03, 2034				
	11564886	Mar 07, 2032	DP U-3531			
	11833166	Apr 03, 2034	U-3776			
	11833166	Apr 03, 2034	U-3777			
	12115179	Feb 11, 2030	U-4023			
	7407955	May 02, 2025	DS DP			
	7407955*PED	Nov 02, 2025				
	7579449	Aug 01, 2028	DS			
	7579449*PED	Feb 01, 2029				
	7713938	Apr 15, 2027	DS DP			
	7713938*PED	Oct 15, 2027				
	8551957	Oct 14, 2029	DP U-2730			
	8551957*PED	Apr 14, 2030				
	8883805	Nov 26, 2025	DP			
	8883805*PED	May 26, 2026				
	9155705	May 21, 2030	DP			
	9415016	Apr 02, 2029	DP			
	9949998	Jun 11, 2034	U-2731			
	9949998*PED	Dec 11, 2034				
<u>EMPAGLIFLOZIN; LINAGLIPTIN; METFORMIN HYDROCHLORIDE - TRIJARDY XR</u>						
N 212614 003	10022379	Apr 02, 2029	DP U-2732			
	10258637	Apr 03, 2034	U-2731			
	10258637*PED	Oct 03, 2034				
	10406172	Jun 15, 2030	DP U-2733			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>EMPAGLIFLOZIN; LINAGLIPTIN; METFORMIN HYDROCHLORIDE - TRIJARDY XR</u>						
N 212614 003	10596120	Mar 07, 2032	DP U-2776			
	10596120	Mar 07, 2032	DP U-2790			
	11090323	Apr 03, 2034	U-3192			
	11090323*PED	Oct 03, 2034				
	11564886	Mar 07, 2032	DP U-3531			
	11833166	Apr 03, 2034	U-3776			
	11833166	Apr 03, 2034	U-3777			
	12115179	Feb 11, 2030	U-4023			
	7407955	May 02, 2025	DS DP			
	7407955*PED	Nov 02, 2025				
	7579449	Aug 01, 2028	DS			
	7579449*PED	Feb 01, 2029				
	7713938	Apr 15, 2027	DS DP			
	7713938*PED	Oct 15, 2027				
	8551957	Oct 14, 2029	DP U-2730			
	8551957*PED	Apr 14, 2030				
	8883805	Nov 26, 2025	DP			
	8883805*PED	May 26, 2026				
	9155705	May 21, 2030	DP			
	9415016	Apr 02, 2029	DP			
	9949998	Jun 11, 2034	U-2731			
	9949998*PED	Dec 11, 2034				
<u>EMPAGLIFLOZIN; LINAGLIPTIN; METFORMIN HYDROCHLORIDE - TRIJARDY XR</u>						
N 212614 004	10022379	Apr 02, 2029	DP U-2732			
	10258637	Apr 03, 2034	U-2731			
	10258637*PED	Oct 03, 2034				
	10406172	Jun 15, 2030	DP U-2733			
	10596120	Mar 07, 2032	DP U-2776			
	10596120	Mar 07, 2032	DP U-2790			
	11090323	Apr 03, 2034	U-3192			
	11090323*PED	Oct 03, 2034				
	11833166	Apr 03, 2034	U-3776			
	11833166	Apr 03, 2034	U-3777			
	12115179	Feb 11, 2030	U-4023			
	7407955	May 02, 2025	DS DP			
	7407955*PED	Nov 02, 2025				
	7579449	Aug 01, 2028	DS			
	7579449*PED	Feb 01, 2029				
	7713938	Apr 15, 2027	DS DP			
	7713938*PED	Oct 15, 2027				
	8551957	Oct 14, 2029	DP U-2730			
	8551957*PED	Apr 14, 2030				
	8883805	Nov 26, 2025	DP			
	8883805*PED	May 26, 2026				
	9155705	May 21, 2030	DP			
	9415016	Apr 02, 2029	DP			
	9949998	Jun 11, 2034	U-2731			
	9949998*PED	Dec 11, 2034				

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>EMPAGLIFLOZIN; METFORMIN HYDROCHLORIDE - SYNJARDY</u>						
N 206111 001	10258637	Apr 03, 2034	U-2290		NPP	Jun 20, 2026
	10258637*PED	Oct 03, 2034			PED	Dec 20, 2026
	10610489	Sep 30, 2030	DP			
	10610489*PED	Mar 30, 2031				
	11090323	Apr 03, 2034	U-3193			
	11090323*PED	Oct 03, 2034				
	11813275	Apr 03, 2034	U-3759			
	11833166	Apr 03, 2034	U-3776			
	11833166	Apr 03, 2034	U-3777			
	12115179	Feb 11, 2030	U-4023			
	7579449	Aug 01, 2028	DS			
	7579449*PED	Feb 01, 2029				
	7713938	Apr 15, 2027	DS DP			
	7713938*PED	Oct 15, 2027				
	9949997	May 17, 2034	U-3532			
	9949997	May 17, 2034	U-3533			
	9949997*PED	Nov 17, 2034				
<u>EMPAGLIFLOZIN; METFORMIN HYDROCHLORIDE - SYNJARDY</u>						
N 206111 002	10258637	Apr 03, 2034	U-2290		NPP	Jun 20, 2026
	10258637*PED	Oct 03, 2034			PED	Dec 20, 2026
	10610489	Sep 30, 2030	DP			
	10610489*PED	Mar 30, 2031				
	11090323	Apr 03, 2034	U-3193			
	11090323*PED	Oct 03, 2034				
	11813275	Apr 03, 2034	U-3759			
	11833166	Apr 03, 2034	U-3776			
	11833166	Apr 03, 2034	U-3777			
	12115179	Feb 11, 2030	U-4023			
	7579449	Aug 01, 2028	DS			
	7579449*PED	Feb 01, 2029				
	7713938	Apr 15, 2027	DS DP			
	7713938*PED	Oct 15, 2027				
	9949997	May 17, 2034	U-3532			
	9949997	May 17, 2034	U-3533			
	9949997*PED	Nov 17, 2034				
<u>EMPAGLIFLOZIN; METFORMIN HYDROCHLORIDE - SYNJARDY</u>						
N 206111 003	10258637	Apr 03, 2034	U-2290		NPP	Jun 20, 2026
	10258637*PED	Oct 03, 2034			PED	Dec 20, 2026
	10610489	Sep 30, 2030	DP			
	10610489*PED	Mar 30, 2031				
	11090323	Apr 03, 2034	U-3193			
	11090323*PED	Oct 03, 2034				
	11833166	Apr 03, 2034	U-3776			
	11833166	Apr 03, 2034	U-3777			
	12115179	Feb 11, 2030	U-4023			
	7579449	Aug 01, 2028	DS			
	7579449*PED	Feb 01, 2029				
	7713938	Apr 15, 2027	DS DP			
	7713938*PED	Oct 15, 2027				

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>EMPAGLIFLOZIN; METFORMIN HYDROCHLORIDE - SYNJARDY</u>						
N 206111 003	9949997	May 17, 2034	U-3532			
	9949997	May 17, 2034	U-3533			
	9949997*PED	Nov 17, 2034				
<u>EMPAGLIFLOZIN; METFORMIN HYDROCHLORIDE - SYNJARDY</u>						
N 206111 004	10258637	Apr 03, 2034	U-2290		NPP	Jun 20, 2026
	10258637*PED	Oct 03, 2034			PED	Dec 20, 2026
	10610489	Sep 30, 2030	DP			
	10610489*PED	Mar 30, 2031				
	11090323	Apr 03, 2034	U-3193			
	11090323*PED	Oct 03, 2034				
	11833166	Apr 03, 2034	U-3776			
	11833166	Apr 03, 2034	U-3777			
	12115179	Feb 11, 2030	U-4023			
	7579449	Aug 01, 2028	DS			
	7579449*PED	Feb 01, 2029				
	7713938	Apr 15, 2027	DS DP			
	7713938*PED	Oct 15, 2027				
	9949997	May 17, 2034	U-3532			
	9949997	May 17, 2034	U-3533			
	9949997*PED	Nov 17, 2034				
<u>EMPAGLIFLOZIN; METFORMIN HYDROCHLORIDE - SYNJARDY XR</u>						
N 208658 001	10258637	Apr 03, 2034	U-2290		M-296	Jun 20, 2026
	10258637*PED	Oct 03, 2034			PED	Dec 20, 2026
	10596120	Mar 07, 2032	DP U-2775			
	10596120	Mar 07, 2032	DP U-2792			
	10596120*PED	Sep 07, 2032				
	11090323	Apr 03, 2034	U-3193			
	11090323*PED	Oct 03, 2034				
	11813275	Apr 03, 2034	U-3759			
	11833166	Apr 03, 2034	U-3776			
	11833166	Apr 03, 2034	U-3777			
	12115179	Feb 11, 2030	U-4023			
	7579449	Aug 01, 2028	DS			
	7579449*PED	Feb 01, 2029				
	7713938	Apr 15, 2027	DS DP			
	7713938*PED	Oct 15, 2027				
	9949997	May 17, 2034	U-3532			
	9949997	May 17, 2034	U-3533			
	9949997*PED	Nov 17, 2034				
	9949998	Jun 11, 2034	U-2290			
	9949998*PED	Dec 11, 2034				
<u>EMPAGLIFLOZIN; METFORMIN HYDROCHLORIDE - SYNJARDY XR</u>						
N 208658 002	10258637	Apr 03, 2034	U-2290		M-296	Jun 20, 2026
	10258637*PED	Oct 03, 2034			PED	Dec 20, 2026
	10596120	Mar 07, 2032	DP U-2775			
	10596120	Mar 07, 2032	DP U-2792			
	10596120*PED	Sep 07, 2032				
	11090323	Apr 03, 2034	U-3193			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>EMPAGLIFLOZIN; METFORMIN HYDROCHLORIDE - SYNJARDY XR</u>						
N 208658 002	11090323*PED	Oct 03, 2034				
	11813275	Apr 03, 2034	U-3759			
	11833166	Apr 03, 2034	U-3776			
	11833166	Apr 03, 2034	U-3777			
	12115179	Feb 11, 2030	U-4023			
	7579449	Aug 01, 2028	DS			
	7579449*PED	Feb 01, 2029				
	7713938	Apr 15, 2027	DS DP			
	7713938*PED	Oct 15, 2027				
	9949997	May 17, 2034	U-3532			
	9949997	May 17, 2034	U-3533			
	9949997*PED	Nov 17, 2034				
	9949998	Jun 11, 2034	U-2290			
	9949998*PED	Dec 11, 2034				
<u>EMPAGLIFLOZIN; METFORMIN HYDROCHLORIDE - SYNJARDY XR</u>						
N 208658 003	10258637	Apr 03, 2034	U-2290		M-296	Jun 20, 2026
	10258637*PED	Oct 03, 2034			PED	Dec 20, 2026
	10596120	Mar 07, 2032	DP U-2775			
	10596120	Mar 07, 2032	DP U-2792			
	10596120*PED	Sep 07, 2032				
	11090323	Apr 03, 2034	U-3193			
	11090323*PED	Oct 03, 2034				
	11833166	Apr 03, 2034	U-3776			
	11833166	Apr 03, 2034	U-3777			
	12115179	Feb 11, 2030	U-4023			
	7579449	Aug 01, 2028	DS			
	7579449*PED	Feb 01, 2029				
	7713938	Apr 15, 2027	DS DP			
	7713938*PED	Oct 15, 2027				
	9949997	May 17, 2034	U-3532			
	9949997	May 17, 2034	U-3533			
	9949997*PED	Nov 17, 2034				
	9949998	Jun 11, 2034	U-2290			
	9949998*PED	Dec 11, 2034				
<u>EMPAGLIFLOZIN; METFORMIN HYDROCHLORIDE - SYNJARDY XR</u>						
N 208658 004	10258637	Apr 03, 2034	U-2290		M-296	Jun 20, 2026
	10258637*PED	Oct 03, 2034			PED	Dec 20, 2026
	10596120	Mar 07, 2032	DP U-2775			
	10596120	Mar 07, 2032	DP U-2792			
	10596120*PED	Sep 07, 2032				
	11090323	Apr 03, 2034	U-3193			
	11090323*PED	Oct 03, 2034				
	11833166	Apr 03, 2034	U-3776			
	11833166	Apr 03, 2034	U-3777			
	12115179	Feb 11, 2030	U-4023			
	7579449	Aug 01, 2028	DS			
	7579449*PED	Feb 01, 2029				
	7713938	Apr 15, 2027	DS DP			
	7713938*PED	Oct 15, 2027				

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>EMPAGLIFLOZIN; METFORMIN HYDROCHLORIDE - SYNJARDY XR</u>						
N 208658	004	9949997	May 17, 2034	U-3532		
		9949997	May 17, 2034	U-3533		
		9949997*PED	Nov 17, 2034			
		9949998	Jun 11, 2034	U-2290		
		9949998*PED	Dec 11, 2034			
<u>EMTRICITABINE; RILPIVIRINE HYDROCHLORIDE; TENOFOVIR ALAFENAMIDE FUMARATE - ODEFSEY</u>						
N 208351	001	7125879	Apr 21, 2025	DS DP U-257		
		7390791	Apr 17, 2025	DS DP		
		7390791*PED	Oct 17, 2025			
		8754065	Aug 15, 2032	DS DP U-257		
		8754065*PED	Feb 15, 2033			
		9296769	Aug 15, 2032	DS DP U-257		
		9296769*PED	Feb 15, 2033			
<u>EMTRICITABINE; RILPIVIRINE HYDROCHLORIDE; TENOFOVIR DISOPROXIL FUMARATE - COMPLERA</u>						
N 202123	001	10857102	Jan 14, 2033	DP		
		7125879	Apr 21, 2025	DS DP U-257		
		8841310	Dec 09, 2025	DP U-257		
<u>EMTRICITABINE; TENOFOVIR ALAFENAMIDE FUMARATE - DESCOVY</u>						
N 208215	001	7390791	Apr 17, 2025	DS DP		
		7390791*PED	Oct 17, 2025			
		8754065	Aug 15, 2032	DS DP U-1259		
		8754065	Aug 15, 2032	DS DP U-1663		
		8754065	Aug 15, 2032	DS DP U-257		
		8754065*PED	Feb 15, 2033			
		9296769	Aug 15, 2032	DS DP U-1259		
		9296769	Aug 15, 2032	DS DP U-1663		
		9296769	Aug 15, 2032	DS DP U-257		
		9296769*PED	Feb 15, 2033			
<u>EMTRICITABINE; TENOFOVIR ALAFENAMIDE FUMARATE - DESCOVY</u>						
N 208215	002	7390791	Apr 17, 2025	DS DP	ODE-457	Jan 07, 2029
		7390791*PED	Oct 17, 2025			
		8754065	Aug 15, 2032	DS DP U-1663		
		8754065*PED	Feb 15, 2033			
		9296769	Aug 15, 2032	DS DP U-1663		
		9296769*PED	Feb 15, 2033			
<u>ENALAPRIL MALEATE - EPANED KIT</u>						
N 204308	001	8568747	Nov 06, 2032	DP		
		8778366	Nov 06, 2032	U-1723		
		8778366	Nov 06, 2032	U-185		
		8778366	Nov 06, 2032	U-1892		
		8778366	Nov 06, 2032	U-3		
		8778366	Nov 06, 2032	U-71		
		9855214	Nov 06, 2032	DP		
		9968553	Nov 06, 2032	U-1723		
		9968553	Nov 06, 2032	U-185		
		9968553	Nov 06, 2032	U-1892		
		9968553	Nov 06, 2032	U-3		
		9968553	Nov 06, 2032	U-71		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>ENALAPRIL MALEATE - EPANED KIT</u>						
N 204308	001 8568747	Nov 06, 2032	DP			
	8778366	Nov 06, 2032	U-1723			
	8778366	Nov 06, 2032	U-185			
	8778366	Nov 06, 2032	U-1892			
	8778366	Nov 06, 2032	U-3			
	8778366	Nov 06, 2032	U-71			
	9855214	Nov 06, 2032	DP			
	9968553	Nov 06, 2032	U-1723			
	9968553	Nov 06, 2032	U-185			
	9968553	Nov 06, 2032	U-1892			
	9968553	Nov 06, 2032	U-3			
	9968553	Nov 06, 2032	U-71			
<u>ENALAPRIL MALEATE - EPANED</u>						
N 208686	001 10039745	Mar 25, 2036	DP			
	10154987	Mar 25, 2036	U-1723			
	10154987	Mar 25, 2036	U-185			
	10154987	Mar 25, 2036	U-1892			
	10154987	Mar 25, 2036	U-3			
	10154987	Mar 25, 2036	U-71			
	10772868	Mar 25, 2036	DP			
	10786482	Mar 25, 2036	DP			
	11040023	Mar 25, 2036	DP			
	11141405	Mar 25, 2036	DP			
	11173141	Mar 25, 2036	DP			
	9669008	Mar 25, 2036	DP			
	9808442	Mar 25, 2036	U-1723			
	9808442	Mar 25, 2036	U-185			
	9808442	Mar 25, 2036	U-1892			
	9808442	Mar 25, 2036	U-3			
	9808442	Mar 25, 2036	U-71			
<u>ENASIDENIB MESYLATE - IDHIFA</u>						
N 209606	001 10093654	Aug 01, 2034	DS DP U-2087			
	10294215	Jan 07, 2033	DP U-2087			
	10610125	Jun 21, 2030	U-2087			
	9512107	Jan 07, 2033	DS DP U-2087			
	9732062	Sep 16, 2034	DS			
	9738625	Aug 01, 2034	DS			
<u>ENASIDENIB MESYLATE - IDHIFA</u>						
N 209606	002 10093654	Aug 01, 2034	DS DP U-2087			
	10294215	Jan 07, 2033	DP U-2087			
	10610125	Jun 21, 2030	U-2087			
	9512107	Jan 07, 2033	DS DP U-2087			
	9732062	Sep 16, 2034	DS			
	9738625	Aug 01, 2034	DS			
<u>ENCORAFENIB - BRAFTOVI</u>						
N 210496	001 10005761	Aug 27, 2030	U-2335		I-928	Oct 11, 2026
	8541575	Feb 26, 2030	DS DP U-2335		ODE-194	Jun 27, 2025
	8946250	Jul 23, 2029	DS DP		ODE-445	Oct 11, 2030

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>ENCORAFENIB - BRAFTOVI</u>						
N 210496 001	9314464	Jul 04, 2031	U-2336			
	9387208	Nov 21, 2032	DP			
	9593099	Aug 27, 2030	DS			
	9593100	Aug 27, 2030	DP			
	9763941	Nov 21, 2032	U-2335			
	9850229	Aug 27, 2030	U-2337			
	9850230	Aug 27, 2030	U-2334			
<u>ENCORAFENIB - BRAFTOVI</u>						
N 210496 002	10005761	Aug 27, 2030	U-2335		I-928	Oct 11, 2026
	10005761	Aug 27, 2030	U-2802		ODE-194	Jun 27, 2025
	10005761	Aug 27, 2030	U-2803		ODE-445	Oct 11, 2030
	10005761	Aug 27, 2030	U-3738			
	10258622	Nov 21, 2032	U-2802			
	8541575	Feb 26, 2030	DS DP U-2335			
	8541575	Feb 26, 2030	DS DP U-2802			
	8541575	Feb 26, 2030	DS DP U-2803			
	8541575	Feb 26, 2030	DS DP U-3738			
	8946250	Jul 23, 2029	DS DP			
	9314464	Jul 04, 2031	U-2336			
	9314464	Jul 04, 2031	U-2802			
	9314464	Jul 04, 2031	U-2803			
	9314464	Jul 04, 2031	U-3738			
	9387208	Nov 21, 2032	DP			
	9474754	Aug 05, 2033	U-2802			
	9593099	Aug 27, 2030	DS			
	9593100	Aug 27, 2030	DP			
	9763941	Nov 21, 2032	U-2335			
	9850229	Aug 27, 2030	U-2337			
	9850230	Aug 27, 2030	U-2334			
	9850230	Aug 27, 2030	U-2802			
	9850230	Aug 27, 2030	U-2803			
	9850230	Aug 27, 2030	U-3738			
	RE49556	Feb 27, 2030	DS DP			
<u>ENSIFENTRINE - OHTUVAYRE</u>						
N 217389 001	10945950	Sep 15, 2035	DP		NCE	Jun 26, 2029
	9062047	Aug 21, 2031	DS U-3962			
	9956171	Sep 15, 2035	DP U-3962			
<u>ENTRECTINIB - ROZLYTREK</u>						
N 212725 001	10231965	Feb 17, 2035	U-2617		NPP	Oct 20, 2026
	10231965	Feb 17, 2035	U-2618		ODE-265	Aug 15, 2026
	10398693	Jul 18, 2038	DP		ODE-313	Aug 15, 2026
	10561651	Feb 19, 2035	U-2745		ODE-448	Oct 20, 2030
	10738037	May 18, 2037	DS DP U-2946			
	11091469	May 18, 2037	U-2617			
	11091469	May 18, 2037	U-2618			
	11253515	Jul 18, 2038	DP			
	8299057	Mar 01, 2029	DS DP			
	8673893	Jul 08, 2028	U-2617			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>ENTRECTINIB - ROZLYTREK</u>						
N 212725 001	8673893	Jul 08, 2028	U-2618			
	9029356	Jul 08, 2028	DS DP			
	9085558	Jul 08, 2028	DP			
	9085565	May 22, 2033	DS DP			
	9255087	Jul 08, 2028	U-2617			
	9255087	Jul 08, 2028	U-2618			
	9616059	Jul 08, 2028	U-2618			
	9649306	May 22, 2033	U-2617			
	9649306	May 22, 2033	U-2618			
<u>ENTRECTINIB - ROZLYTREK</u>						
N 212725 002	10231965	Feb 17, 2035	U-2617		NPP	Oct 20, 2026
	10231965	Feb 17, 2035	U-2618		ODE-265	Aug 15, 2026
	10398693	Jul 18, 2038	DP		ODE-313	Aug 15, 2026
	10561651	Feb 19, 2035	U-2745		ODE-448	Oct 20, 2030
	10738037	May 18, 2037	DS DP U-2946			
	11091469	May 18, 2037	U-2617			
	11091469	May 18, 2037	U-2618			
	11253515	Jul 18, 2038	DP			
	8299057	Mar 01, 2029	DS DP			
	8673893	Jul 08, 2028	U-2617			
	8673893	Jul 08, 2028	U-2618			
	9029356	Jul 08, 2028	DS DP			
	9085558	Jul 08, 2028	DP			
	9085565	May 22, 2033	DS DP			
	9255087	Jul 08, 2028	U-2617			
	9255087	Jul 08, 2028	U-2618			
	9616059	Jul 08, 2028	U-2618			
	9649306	May 22, 2033	U-2617			
	9649306	May 22, 2033	U-2618			
<u>ENTRECTINIB - ROZLYTREK</u>						
N 218550 001					NP	Oct 20, 2026
					ODE-448	Oct 20, 2030
<u>ENZALUTAMIDE - XTANDI</u>						
N 203415 001	7709517	Aug 13, 2027	DS DP		I-926	Nov 17, 2026
	8183274	Aug 24, 2026	U-1281			
	8183274	Aug 24, 2026	U-1588			
	8183274	Aug 24, 2026	U-2345			
	8183274	Aug 24, 2026	U-2708			
	8183274	Aug 24, 2026	U-3763			
	9126941	May 15, 2026	U-1588			
	9126941	May 15, 2026	U-2345			
	9126941	May 15, 2026	U-2708			
	9126941	May 15, 2026	U-3763			
<u>ENZALUTAMIDE - XTANDI</u>						
N 213674 001	11839689	Sep 11, 2033	DP		I-926	Nov 17, 2026
	7709517	Aug 13, 2027	DS DP			
	8183274	Aug 24, 2026	U-2345			
	8183274	Aug 24, 2026	U-2708			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>ENZALUTAMIDE - XTANDI</u>						
N 213674	001	8183274	Aug 24, 2026	U-3763		
		9126941	May 15, 2026	U-2345		
		9126941	May 15, 2026	U-2708		
		9126941	May 15, 2026	U-3763		
<u>ENZALUTAMIDE - XTANDI</u>						
N 213674	002	11839689	Sep 11, 2033	DP	I-926	Nov 17, 2026
		7709517	Aug 13, 2027	DS DP		
		8183274	Aug 24, 2026	U-2345		
		8183274	Aug 24, 2026	U-2708		
		8183274	Aug 24, 2026	U-3763		
		9126941	May 15, 2026	U-2345		
		9126941	May 15, 2026	U-2708		
		9126941	May 15, 2026	U-3763		
<u>EPHEDRINE SULFATE - EPHEDRINE SULFATE</u>						
A 217721	001				CGT	Jun 18, 2025
<u>EPHEDRINE SULFATE - EMERPHED</u>						
N 213407	001	11090278	May 16, 2040	U-3183		
		11241400	May 16, 2040	U-3183		
		11464752	May 16, 2040	DP		
		11478436	May 16, 2040	U-3183		
		11571398	May 16, 2040	U-3183		
<u>EPHEDRINE SULFATE - EMERPHED</u>						
N 213407	002	11464752	May 16, 2040	DP		
		11571398	May 16, 2040	U-3183		
<u>EPHEDRINE SULFATE - EPHEDRINE SULFATE</u>						
N 213994	001	10869845	Jan 22, 2040	DP		
<u>EPHEDRINE SULFATE - EPHEDRINE SULFATE</u>						
N 213994	002	10869845	Jan 22, 2040	DP		
		12029710	Jan 22, 2040	U-3963		
<u>EPINEPHRINE - EPIPEN</u>						
N 019430	001	7449012	Sep 11, 2025	DP		
		7794432	Sep 11, 2025	DP		
		8048035	Sep 11, 2025	DP		
		8870827	Sep 11, 2025	DP		
		9586010	Sep 11, 2025	DP		
<u>EPINEPHRINE - EPIPEN JR.</u>						
N 019430	002	7449012	Sep 11, 2025	DP		
		7794432	Sep 11, 2025	DP		
		8048035	Sep 11, 2025	DP		
		8870827	Sep 11, 2025	DP		
		9586010	Sep 11, 2025	DP		
<u>EPINEPHRINE - AUVI-Q</u>						
N 201739	001	10335549	Apr 30, 2025	DP		
		11590286	Dec 12, 2026	DP		
		7947017	Mar 12, 2028	DP		
		8021344	Nov 02, 2029	DP		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>EPINEPHRINE - AUVI-Q</u>						
N 201739	001	8206360	Feb 27, 2027	DP		
		8226610	Apr 10, 2029	DP		
		8231573	Nov 25, 2028	DP		
<u>EPINEPHRINE - AUVI-Q</u>						
N 201739	002	10335549	Apr 30, 2025	DP		
		10688244	Dec 21, 2037	DP U-2980		
		10842938	Dec 21, 2037	DP U-2980		
		11590286	Dec 12, 2026	DP		
		11771830	Dec 21, 2037	DP U-2980		
		7947017	Mar 12, 2028	DP		
		8021344	Nov 02, 2029	DP		
		8206360	Feb 27, 2027	DP		
		8226610	Apr 10, 2029	DP		
		8231573	Nov 25, 2028	DP		
<u>EPINEPHRINE - AUVI-Q</u>						
N 201739	003	10335549	Apr 30, 2025	DP		
		10688244	Dec 21, 2037	DP U-2980		
		10842938	Dec 21, 2037	DP U-2980		
		11590286	Dec 12, 2026	DP		
		11771830	Dec 21, 2037	DP U-2980		
		7947017	Mar 12, 2028	DP		
		8021344	Nov 02, 2029	DP		
		8206360	Feb 27, 2027	DP		
		8226610	Apr 10, 2029	DP		
		8231573	Nov 25, 2028	DP		
<u>EPINEPHRINE - ADRENALIN</u>						
N 204200	001	9119876	Mar 13, 2035	DP		
		9295657	Mar 13, 2035		U-1829	
<u>EPINEPHRINE - ADRENALIN</u>						
N 204640	001	10130592	Mar 13, 2035	DP		
		9119876	Mar 13, 2035	DP		
		9295657	Mar 13, 2035		U-1829	
<u>EPINEPHRINE - EPINEPHRINE</u>						
N 205029	001	10004700	Aug 14, 2034	DP U-2325		Y
		10039728	Aug 14, 2034		U-1828	Y
		9283197	Aug 15, 2034	DP U-1828		Y
		9283197	Aug 15, 2034	DP U-1829		Y
		9283197	Aug 15, 2034	DP U-1830		Y
<u>EPINEPHRINE - PRIMATENE MIST</u>						
N 205920	001	8367734	Jan 26, 2026	DP		
<u>EPINEPHRINE - SYMJEPI</u>						
N 207534	001	11141540	Oct 20, 2036	DP U-3379		
<u>EPINEPHRINE - SYMJEPI</u>						
N 207534	002	11141540	Oct 20, 2036	DP U-3379		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>EPINEPHRINE - NEFFY</u>						
N 214697	001	10576156	Feb 06, 2038	DP		
		10682414	Feb 06, 2039	U-3979		
		11173209	Feb 06, 2038	U-3979		
		11191838	Feb 06, 2039	DP U-3979		
		11717571	Feb 06, 2039	U-3979		
		11744895	Feb 06, 2039	U-3979		
		11918655	Feb 06, 2039	DP U-3979		
<u>EPINEPHRINE - ADRENALIN</u>						
N 215875	001	10653646	Mar 21, 2039	DP		
		11083698	Mar 21, 2039	U-3567		
		11207280	Mar 21, 2039	DP		
		12133837	Mar 21, 2039	DP		
<u>EPINEPHRINE - ADRENALIN</u>						
N 215875	002	10653646	Mar 21, 2039	DP		
		11083698	Mar 21, 2039	U-3567		
		11207280	Mar 21, 2039	DP		
		12133837	Mar 21, 2039	DP		
<u>EPINEPHRINE - ADRENALIN</u>						
N 215875	003	10653646	Mar 21, 2039	DP		
		11083698	Mar 21, 2039	U-3567		
		11207280	Mar 21, 2039	DP		
		12133837	Mar 21, 2039	DP		
<u>EPINEPHRINE - ADRENALIN</u>						
N 215875	004	10653646	Mar 21, 2039	DP		
		11083698	Mar 21, 2039	U-3567		
		11207280	Mar 21, 2039	DP		
		12133837	Mar 21, 2039	DP		
<u>EPINEPHRINE - ADRENALIN</u>						
N 215875	005	10653646	Mar 21, 2039	DP		
		11083698	Mar 21, 2039	U-3567		
		11207280	Mar 21, 2039	DP		
		12133837	Mar 21, 2039	DP		
<u>EPLONTERSEN SODIUM - WAINUA (AUTOINJECTOR)</u>						
N 217388	001	10683499	Aug 25, 2034	DS DP U-2378	NCE	Dec 21, 2028
		8101743	Apr 01, 2025	DS DP	ODE-461	Dec 21, 2030
		9127276	May 01, 2034	DS		
		9181549	May 01, 2034	DS		
<u>EPOPROSTENOL SODIUM - VELETRI</u>						
N 022260	001	8318802	Mar 15, 2027	DP		
		8598227	Feb 02, 2027			
<u>EPOPROSTENOL SODIUM - VELETRI</u>						
N 022260	002	8318802	Mar 15, 2027	DP		
		8598227	Feb 02, 2027			
<u>EPAVACYCLINE DIHYDROCHLORIDE - XERAVA</u>						
N 211109	001	10961190	Oct 19, 2037	DS	NCE	Aug 27, 2023
		11578044	Oct 19, 2037	DS	GAIN	Aug 27, 2028

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>ERAVACYCLINE DIHYDROCHLORIDE - XERAVA</u>						
N 211109	001 8796245	Aug 07, 2029	U-2380			
	8906887	Dec 28, 2030	DP			
<u>ERAVACYCLINE DIHYDROCHLORIDE - XERAVA</u>						
N 211109	002 10961190	Oct 19, 2037	DS		NCE	Aug 27, 2023
	11578044	Oct 19, 2037	DS		GAIN	Aug 27, 2028
	8796245	Aug 07, 2029	U-2380			
	8906887	Dec 28, 2030	DP			
<u>ERDAFITINIB - BALVERSA</u>						
N 212018	001 10478494	Aug 13, 2036	U-3902		I-930	Jan 19, 2027
	10898482	Feb 09, 2036	DP U-2518			
	10898482	Feb 09, 2036	DP U-3065			
	10898482	Feb 09, 2036	DP U-3066			
	10898482	Feb 09, 2036	DP U-3067			
	10898482	Feb 09, 2036	DP U-3805			
	10898482	Feb 09, 2036	DP U-3806			
	10898482	Feb 09, 2036	DP U-3807			
	11077106	Feb 02, 2038	U-3196			
	11077106	Feb 02, 2038	U-3808			
	11684620	Feb 09, 2036	U-3805			
	11684620	Feb 09, 2036	U-3806			
	11684620	Feb 09, 2036	U-3807			
	12037644	Oct 18, 2035	U-3972			
	8895601	Apr 12, 2033	DS DP			
	9464071	Apr 28, 2031	U-3805			
	9902714	Mar 26, 2035	DP			
<u>ERDAFITINIB - BALVERSA</u>						
N 212018	002 10478494	Aug 13, 2036	U-3902		I-930	Jan 19, 2027
	10898482	Feb 09, 2036	DP U-2518			
	10898482	Feb 09, 2036	DP U-3065			
	10898482	Feb 09, 2036	DP U-3066			
	10898482	Feb 09, 2036	DP U-3067			
	10898482	Feb 09, 2036	DP U-3805			
	10898482	Feb 09, 2036	DP U-3806			
	10898482	Feb 09, 2036	DP U-3807			
	11077106	Feb 02, 2038	U-3196			
	11077106	Feb 02, 2038	U-3808			
	11684620	Feb 09, 2036	U-3805			
	11684620	Feb 09, 2036	U-3806			
	11684620	Feb 09, 2036	U-3807			
	12037644	Oct 18, 2035	U-3972			
	8895601	Apr 12, 2033	DS DP			
	9464071	Apr 28, 2031	U-3805			
	9902714	Mar 26, 2035	DP			
<u>ERDAFITINIB - BALVERSA</u>						
N 212018	003 10478494	Aug 13, 2036	U-3902		I-930	Jan 19, 2027
	10898482	Feb 09, 2036	DP U-2518			
	10898482	Feb 09, 2036	DP U-3065			
	10898482	Feb 09, 2036	DP U-3066			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>ERDAFITINIB - BALVERSA</u>						
N 212018	003 10898482	Feb 09, 2036	DP U-3067			
	10898482	Feb 09, 2036	DP U-3805			
	10898482	Feb 09, 2036	DP U-3806			
	10898482	Feb 09, 2036	DP U-3807			
	11077106	Feb 02, 2038	U-3196			
	11077106	Feb 02, 2038	U-3808			
	11684620	Feb 09, 2036	U-3805			
	11684620	Feb 09, 2036	U-3806			
	11684620	Feb 09, 2036	U-3807			
	12037644	Oct 18, 2035	U-3972			
	8895601	Apr 12, 2033	DS DP			
	9464071	Apr 28, 2031	U-3805			
	9902714	Mar 26, 2035	DP			
<u>ERIBULIN MESYLATE - HALAVEN</u>						
N 201532	001 RE46965	Jan 08, 2027	DP		M-280	Sep 13, 2025
	RE46965*PED	Jul 08, 2027			PED	Mar 13, 2026
<u>ERTUGLIFLOZIN - STEGLATRO</u>						
N 209803	001 8080580	Jul 13, 2030	DS DP U-2214			
<u>ERTUGLIFLOZIN - STEGLATRO</u>						
N 209803	002 8080580	Jul 13, 2030	DS DP U-2214			
<u>ERTUGLIFLOZIN; METFORMIN HYDROCHLORIDE - SEGLUOMET</u>						
N 209806	001 8080580	Jul 13, 2030	DS DP U-2214			
	9308204	Oct 21, 2030	DP			
	9439902	Oct 21, 2030	U-2214			
<u>ERTUGLIFLOZIN; METFORMIN HYDROCHLORIDE - SEGLUOMET</u>						
N 209806	002 8080580	Jul 13, 2030	DS DP U-2214			
	9308204	Oct 21, 2030	DP			
	9439902	Oct 21, 2030	U-2214			
<u>ERTUGLIFLOZIN; METFORMIN HYDROCHLORIDE - SEGLUOMET</u>						
N 209806	003 8080580	Jul 13, 2030	DS DP U-2214			
	9308204	Oct 21, 2030	DP			
	9439902	Oct 21, 2030	U-2214			
<u>ERTUGLIFLOZIN; METFORMIN HYDROCHLORIDE - SEGLUOMET</u>						
N 209806	004 8080580	Jul 13, 2030	DS DP U-2214			
	9308204	Oct 21, 2030	DP			
	9439902	Oct 21, 2030	U-2214			
<u>ERTUGLIFLOZIN; SITAGLIPTIN PHOSPHATE - STEGLUJAN</u>						
N 209805	001 7326708	Nov 24, 2026	DS DP U-2214			
	7326708*PED	May 24, 2027				
	8080580	Jul 13, 2030	DS DP U-2214			
	9308204	Oct 21, 2030	DP			
	9439901	Oct 21, 2030	U-2214			
<u>ERTUGLIFLOZIN; SITAGLIPTIN PHOSPHATE - STEGLUJAN</u>						
N 209805	002 7326708	Nov 24, 2026	DS DP U-2214			
	7326708*PED	May 24, 2027				
	8080580	Jul 13, 2030	DS DP U-2214			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>ERTUGLIFLOZIN; SITAGLIPTIN PHOSPHATE - STEGLUJAN</u>						
N 209805 002	9308204	Oct 21, 2030	DP			
	9439901	Oct 21, 2030	U-2214			
<u>ERYTHROMYCIN - ERYTHROMYCIN</u>						
A 211975 003					CGT	Apr 22, 2025
<u>ESCITALOPRAM OXALATE - LEXAPRO</u>						
N 021323 001					NPP	May 12, 2026
<u>ESCITALOPRAM OXALATE - LEXAPRO</u>						
N 021323 002					NPP	May 12, 2026
<u>ESCITALOPRAM OXALATE - LEXAPRO</u>						
N 021323 003					NPP	May 12, 2026
<u>ESCITALOPRAM OXALATE - LEXAPRO</u>						
N 021365 001					NPP	May 12, 2026
<u>ESKETAMINE HYDROCHLORIDE - SPRAVATO</u>						
N 211243 001	10869844	Sep 10, 2035	U-3034			
	10869844	Sep 10, 2035	U-3035			
	10869844	Sep 10, 2035	U-3036			
	11173134	Sep 10, 2035	U-3257			
	11173134	Sep 10, 2035	U-3536			
	11311500	Sep 10, 2035	U-3034			
	11311500	Sep 10, 2035	U-3035			
	11311500	Sep 10, 2035	U-3036			
	11446260	Mar 14, 2034	U-3444			
	11446260	Mar 14, 2034	U-3445			
	11446260	Mar 14, 2034	U-3446			
	11883526	Feb 18, 2040	U-3812			
	11883526	Feb 18, 2040	U-3813			
	8785500	Mar 05, 2033	U-2502			
	9592207	Mar 20, 2027	U-2502			
<u>ESLICARBAZEPINE ACETATE - APTIOM</u>						
N 022416 001	10675287	May 06, 2025	U-2041			
	10675287	May 06, 2025	U-2831			
	10695354	May 06, 2025	U-2501			
	10695354	May 06, 2025	U-2831			
	10702536	May 06, 2025	U-2501			
	10912781	Oct 23, 2028	DP			
	11364247	May 06, 2025	U-2501			
	11364247	May 06, 2025	U-2831			
	8372431	Apr 17, 2030	DP			
	9206135	Apr 21, 2026	DS			
	9566244	Oct 23, 2028	DP			
	9643929	Apr 21, 2026	DP			
	9750747	Aug 24, 2032	U-2041			
	9750747	Aug 24, 2032	U-2121			
	9763954	Sep 13, 2028	U-2123			
<u>ESLICARBAZEPINE ACETATE - APTIOM</u>						
N 022416 002	10675287	May 06, 2025	U-2041			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>ESLICARBAZEPINE ACETATE - APTIOM</u>						
N 022416 002	10675287	May 06, 2025	U-2831			
	10695354	May 06, 2025	U-2501			
	10695354	May 06, 2025	U-2831			
	10702536	May 06, 2025	U-2501			
	10912781	Oct 23, 2028	DP			
	11364247	May 06, 2025	U-2501			
	11364247	May 06, 2025	U-2831			
	8372431	Apr 17, 2030	DP			
	9206135	Apr 21, 2026	DS			
	9566244	Oct 23, 2028	DP			
	9643929	Apr 21, 2026	DP			
	9750747	Aug 24, 2032	U-2041			
	9750747	Aug 24, 2032	U-2121			
	9763954	Sep 13, 2028	U-2123			
<u>ESLICARBAZEPINE ACETATE - APTIOM</u>						
N 022416 003	10675287	May 06, 2025	U-2041			
	10675287	May 06, 2025	U-2831			
	10695354	May 06, 2025	U-2501			
	10695354	May 06, 2025	U-2831			
	10702536	May 06, 2025	U-2501			
	10912781	Oct 23, 2028	DP			
	11364247	May 06, 2025	U-2501			
	11364247	May 06, 2025	U-2831			
	8372431	Apr 17, 2030	DP			
	9206135	Apr 21, 2026	DS			
	9566244	Oct 23, 2028	DP			
	9643929	Apr 21, 2026	DP			
	9750747	Aug 24, 2032	U-2041			
	9750747	Aug 24, 2032	U-2121			
	9763954	Sep 13, 2028	U-2123			
<u>ESLICARBAZEPINE ACETATE - APTIOM</u>						
N 022416 004	10675287	May 06, 2025	U-2041			
	10675287	May 06, 2025	U-2831			
	10695354	May 06, 2025	U-2501			
	10695354	May 06, 2025	U-2831			
	10702536	May 06, 2025	U-2501			
	10912781	Oct 23, 2028	DP			
	11364247	May 06, 2025	U-2501			
	11364247	May 06, 2025	U-2831			
	8372431	Apr 17, 2030	DP			
	9206135	Apr 21, 2026	DS			
	9566244	Oct 23, 2028	DP			
	9643929	Apr 21, 2026	DP			
	9750747	Aug 24, 2032	U-2041			
	9750747	Aug 24, 2032	U-2121			
	9763954	Sep 13, 2028	U-2123			
<u>ESMOLOL HYDROCHLORIDE - ESMOLOL HYDROCHLORIDE IN PLASTIC CONTAINER</u>						
N 205703 001	8829054	Mar 15, 2033	DP			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>ESMOLOL HYDROCHLORIDE - ESMOLOL HYDROCHLORIDE IN PLASTIC CONTAINER</u>						
N 205703	001 8835505	Mar 15, 2033	DP			
<u>ESMOLOL HYDROCHLORIDE - ESMOLOL HYDROCHLORIDE DOUBLE STRENGTH IN PLASTIC CONTAINER</u>						
N 205703	002 8829054	Mar 15, 2033	DP			
	8835505	Mar 15, 2033	DP			
<u>ESOMEPRAZOLE MAGNESIUM - ESOMEPRAZOLE MAGNESIUM</u>						
N 214278	001 10076494	Dec 08, 2036	DP			
	10835488	Dec 08, 2036	DP			
<u>ESTRADIOL - MINIVELLE</u>						
N 203752	001 8231906	Jul 04, 2030	DS DP			
	9730900	Jul 10, 2028			U-2086	
	9833419	Jul 10, 2028	DP			
<u>ESTRADIOL - MINIVELLE</u>						
N 203752	002 8231906	Jul 04, 2030	DS DP			
	9730900	Jul 10, 2028			U-2086	
	9833419	Jul 10, 2028	DP			
<u>ESTRADIOL - MINIVELLE</u>						
N 203752	003 8231906	Jul 04, 2030	DS DP			
	9730900	Jul 10, 2028			U-2086	
	9833419	Jul 10, 2028	DP			
<u>ESTRADIOL - MINIVELLE</u>						
N 203752	004 8231906	Jul 04, 2030	DS DP			
	9730900	Jul 10, 2028			U-2086	
	9833419	Jul 10, 2028	DP			
<u>ESTRADIOL - MINIVELLE</u>						
N 203752	005 8231906	Jul 04, 2030	DS DP			
	9724310	Jul 10, 2028	DS DP			
	9730900	Jul 10, 2028	DP		U-2086	
	9833419	Jul 10, 2028	DP			
<u>ESTRADIOL - IMVEXXY</u>						
N 208564	001 10258630	Nov 21, 2032			U-2316	
	10258630	Nov 21, 2032			U-2317	
	10398708	Nov 21, 2032			U-2317	
	10398708	Nov 21, 2032			U-2614	
	10471072	Nov 21, 2032			U-2316	
	10471072	Nov 21, 2032			U-2317	
	10537581	Nov 21, 2032	DP		U-2316	
	10537581	Nov 21, 2032	DP		U-2317	
	10568891	Nov 21, 2032			U-2316	
	10568891	Nov 21, 2032			U-2317	
	10668082	Nov 21, 2032			U-2316	
	10668082	Nov 21, 2032			U-2317	
	10806697	Nov 21, 2032	DP			
	10835487	Nov 21, 2032			U-2316	
	10835487	Nov 21, 2032			U-2317	
	10888516	Nov 21, 2032			U-2316	
	10888516	Nov 21, 2032			U-2317	

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>ESTRADIOL - IMVEXXY</u>						
N 208564 001	11065197	Nov 21, 2032	DP			
	11116717	Nov 21, 2032	DP			
	11123283	Nov 21, 2032	DP			
	11241445	Nov 21, 2032	U-2316			
	11241445	Nov 21, 2032	U-2317			
	11246875	Nov 21, 2032	U-2316			
	11246875	Nov 21, 2032	U-2317			
	11266661	Feb 02, 2034	U-2316			
	11266661	Feb 02, 2034	U-2317			
	11304959	Nov 21, 2032	DP			
	11351182	Nov 21, 2032	U-2316			
	11351182	Nov 21, 2032	U-2317			
	9180091	Nov 21, 2032	DP U-2316			
	9180091	Nov 21, 2032	DP U-2317			
	9289382	Nov 21, 2032	DP			
<u>ESTRADIOL - IMVEXXY</u>						
N 208564 002	10258630	Nov 21, 2032	U-2316			
	10258630	Nov 21, 2032	U-2317			
	10398708	Nov 21, 2032	U-2317			
	10398708	Nov 21, 2032	U-2614			
	10471072	Nov 21, 2032	U-2316			
	10471072	Nov 21, 2032	U-2317			
	10537581	Nov 21, 2032	DP U-2316			
	10537581	Nov 21, 2032	DP U-2317			
	10568891	Nov 21, 2032	U-2316			
	10568891	Nov 21, 2032	U-2317			
	10668082	Nov 21, 2032	U-2316			
	10668082	Nov 21, 2032	U-2317			
	10806697	Nov 21, 2032	DP			
	10835487	Nov 21, 2032	U-2316			
	10835487	Nov 21, 2032	U-2317			
	10888516	Nov 21, 2032	U-2316			
	10888516	Nov 21, 2032	U-2317			
	11065197	Nov 21, 2032	DP			
	11116717	Nov 21, 2032	DP			
	11123283	Nov 21, 2032	DP			
	11241445	Nov 21, 2032	U-2316			
	11241445	Nov 21, 2032	U-2317			
	11246875	Nov 21, 2032	U-2316			
	11246875	Nov 21, 2032	U-2317			
	11266661	Feb 02, 2034	U-2316			
	11266661	Feb 02, 2034	U-2317			
	11304959	Nov 21, 2032	DP			
	11351182	Nov 21, 2032	U-2316			
	11351182	Nov 21, 2032	U-2317			
	11497709	Nov 21, 2032	U-2316			
	11497709	Nov 21, 2032	U-2317			
	9180091	Nov 21, 2032	DP U-2316			
	9180091	Nov 21, 2032	DP U-2317			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>ESTRADIOL - IMVEXXY</u>						
N 208564	002 9289382	Nov 21, 2032	DP			
<u>ESTRADIOL ACETATE - FEMTRACE</u>						
N 021633	001 7572779	Oct 02, 2025	U-904			
<u>ESTRADIOL ACETATE - FEMTRACE</u>						
N 021633	002 7572779	Oct 02, 2025	U-904			
<u>ESTRADIOL ACETATE - FEMTRACE</u>						
N 021633	003 7572779	Oct 02, 2025	U-904			
<u>ESTRADIOL; NORETHINDRONE ACETATE; RELUGOLIX - MYFEMBREE</u>						
N 214846	001 11033551	Sep 29, 2037	U-3129		I-898	Aug 05, 2025
	11793812	May 03, 2038	U-2360		M-289	Jan 27, 2026
	11795178	Sep 27, 2033	DS DP		NCE	Dec 18, 2025
	11957684	Sep 29, 2037	U-3129			
	7300935	Jan 28, 2026	DS			
	8058280	Jan 28, 2026	DS DP			
<u>ESTRADIOL; PROGESTERONE - BIJUVA</u>						
N 210132	001 10052386	Nov 21, 2032	DP			
	10206932	Nov 21, 2032	U-2439			
	10639375	Nov 21, 2032	DP			
	10675288	Nov 21, 2032	U-2439			
	10806740	Nov 21, 2032	DP U-2439			
	11033626	Nov 21, 2032	DP U-2439			
	11103513	Nov 21, 2032	U-2439			
	11103516	Nov 21, 2032	DP			
	11110099	Nov 21, 2032	DP			
	11529360	Nov 21, 2032	DP			
	11793819	Nov 21, 2032	U-2439			
	11865179	Nov 21, 2032	DS DP			
	8633178	Nov 21, 2032	DP			
	8846648	Nov 21, 2032	U-2439			
	8846649	Nov 21, 2032	DP U-2439			
	8987237	Nov 21, 2032	DP			
	8993548	Nov 21, 2032	DP			
	8993549	Nov 21, 2032	DP			
	9006222	Nov 21, 2032	DP U-2439			
	9114145	Nov 21, 2032	U-2439			
	9114146	Nov 21, 2032	DP U-2439			
	9301920	Nov 21, 2032	DP U-2439			
<u>ESTRADIOL; PROGESTERONE - BIJUVA</u>						
N 210132	002 10052386	Nov 21, 2032	DP			
	10206932	Nov 21, 2032	U-2439			
	10675288	Nov 21, 2032	U-2439			
	11033626	Nov 21, 2032	DP U-2439			
	11166963	Nov 21, 2032	DP			
	11793819	Nov 21, 2032	U-2439			
	11865179	Nov 21, 2032	DS DP			
	8633178	Nov 21, 2032	DP			
	8846648	Nov 21, 2032	U-2439			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>ESTRADIOL; PROGESTERONE - BIJUVA</u>						
N 210132	002 8846649	Nov 21, 2032	DP U-2439			
	8933059	Nov 21, 2032	DP U-2439			
	8987237	Nov 21, 2032	DP			
	8993548	Nov 21, 2032	DP			
	8993549	Nov 21, 2032	DP			
	9114145	Nov 21, 2032	U-2439			
	9114146	Nov 21, 2032	DP U-2439			
	9301920	Nov 21, 2032	DP U-2439			
<u>ETELALCETIDE - PARSABIV</u>						
N 208325	001 10344765	Jun 27, 2034	DP			
	11162500	Jun 27, 2034	DP			
	11959486	Jun 27, 2034	DP U-2014			
	8377880	Jul 29, 2030	DS DP			
	8999932	Feb 07, 2031	DS DP U-2014			
	9278995	Jul 29, 2030	DS			
	9701712	Jul 29, 2030	DS DP U-2014			
	9820938	Jun 27, 2034	DP			
<u>ETELALCETIDE - PARSABIV</u>						
N 208325	002 10344765	Jun 27, 2034	DP			
	11162500	Jun 27, 2034	DP			
	11959486	Jun 27, 2034	DP U-2014			
	8377880	Jul 29, 2030	DS DP			
	8999932	Feb 07, 2031	DS DP U-2014			
	9278995	Jul 29, 2030	DS			
	9701712	Jul 29, 2030	DS DP U-2014			
	9820938	Jun 27, 2034	DP			
<u>ETELALCETIDE - PARSABIV</u>						
N 208325	003 10344765	Jun 27, 2034	DP			
	11162500	Jun 27, 2034	DP			
	11959486	Jun 27, 2034	DP U-2014			
	8377880	Jul 29, 2030	DS DP			
	8999932	Feb 07, 2031	DS DP U-2014			
	9278995	Jul 29, 2030	DS			
	9701712	Jul 29, 2030	DS DP U-2014			
	9820938	Jun 27, 2034	DP			
<u>ETEPLIRSEN - EXONDYS 51</u>						
N 206488	001 10337003	Mar 14, 2034	U-1918			
	10364431	Mar 14, 2034	U-1918			
	10364431	Mar 14, 2034	U-1919			
	10781451	Jun 28, 2025	DS DP			
	9018368	Jun 28, 2025	DS DP			
	9243245	Oct 27, 2028	DS U-2097			
	9243245	Oct 27, 2028	DS U-2098			
	9506058	Mar 14, 2034	U-1918			
	9506058	Mar 14, 2034	U-1919			
	RE47751	Jun 28, 2025	U-1918			
	RE47751	Jun 28, 2025	U-2664			
	RE47751	Jun 28, 2025	U-2673			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>ETEPLIRSEN - EXONDYS 51</u>						
N 206488	001	RE47751	Jun 28, 2025		U-2674	
		RE47769	Feb 02, 2029		DP	
		RE48468	Oct 27, 2028		U-2097	
<u>ETEPLIRSEN - EXONDYS 51</u>						
N 206488	002	10337003	Mar 14, 2034		U-1918	
		10364431	Mar 14, 2034		U-1918	
		10364431	Mar 14, 2034		U-1919	
		10781451	Jun 28, 2025	DS DP		
		9018368	Jun 28, 2025	DS DP		
		9243245	Oct 27, 2028	DS	U-2097	
		9243245	Oct 27, 2028	DS	U-2098	
		9506058	Mar 14, 2034		U-1918	
		9506058	Mar 14, 2034		U-1919	
		RE47751	Jun 28, 2025		U-1918	
		RE47751	Jun 28, 2025		U-2664	
		RE47751	Jun 28, 2025		U-2673	
		RE47751	Jun 28, 2025		U-2674	
		RE47769	Feb 02, 2029		DP	
		RE48468	Oct 27, 2028		U-2097	
<u>ETHINYL ESTRADIOL; ETHINYL ESTRADIOL; LEVONORGESTREL - SEASONIQUE</u>						
N 021840	001	7855190	Dec 05, 2028		U-1	
<u>ETHINYL ESTRADIOL; LEVONORGESTREL - LOSEASONIQUE</u>						
N 022262	001	7855190	Dec 05, 2028		U-1	
<u>ETHINYL ESTRADIOL; LEVONORGESTREL - TWIRLA</u>						
N 204017	001	8246978	Aug 26, 2028		DP	
		8747888	Jul 10, 2028		DP	
		9050348	Jul 10, 2028		DP	
<u>ETHINYL ESTRADIOL; LEVONORGESTREL - QUARTETTE</u>						
N 204061	001	8415332	Mar 11, 2029		DP	
		8450299	Oct 07, 2025		U-1	
<u>ETHINYL ESTRADIOL; LEVONORGESTREL - BALCOLTRA</u>						
N 208612	001	7838042	Jun 01, 2027	DS	U-3251	
<u>ETHINYL ESTRADIOL; NORETHINDRONE ACETATE - LO LOESTRIN FE</u>						
N 022501	001	7704984	Feb 02, 2029		U-1090	
<u>ETHINYL ESTRADIOL; NORETHINDRONE ACETATE - LO MINASTRIN FE</u>						
N 204654	001	7704984	Feb 02, 2029		U-1	
<u>ETHINYL ESTRADIOL; NORETHINDRONE ACETATE - FEMLYV</u>						
N 218718	001				NP	Jul 22, 2027
<u>ETHINYL ESTRADIOL; SEGESTERONE ACETATE - ANNOVERA</u>						
N 209627	001	10632066	Feb 01, 2039		U-2786	
		10632066	Feb 01, 2039		U-2787	
		10765628	Feb 01, 2039		U-2786	
		10765628	Feb 01, 2039		U-2787	
		10780047	Feb 01, 2039		U-2786	
		10780047	Feb 01, 2039		U-2787	
		10918649	Jun 21, 2039		DP	

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>ETHINYL ESTRADIOL; SEGESTERONE ACETATE - ANNOVERA</u>						
N 209627	001	10925882	Jun 21, 2039	DP		
		10940157	Jun 21, 2039	DP		
		11529308	Jun 21, 2039	DP		
		11850251	Jun 21, 2039	U-3785		
<u>ETONOGESTREL - IMPLANON</u>						
N 021529	001	9757552	Jul 28, 2030	DP U-1		
<u>ETONOGESTREL - NEXPLANON</u>						
N 021529	002	10821277	May 31, 2027	DP		
		8722037	Sep 28, 2027	DP		
		8888745	Aug 28, 2026	DP		
		9757552	Jul 28, 2030	DP U-1		
<u>ETRASIMOD ARGININE - VELSIPITY</u>						
N 216956	001	10301262	Jun 21, 2036	DS DP	NCE	Oct 12, 2028
		10676435	Jun 21, 2036	U-3731		
		11007175	Jan 06, 2036	U-3730		
		11091435	Jun 21, 2036	DS DP		
		11884626	Jun 21, 2036	U-3731		
		12156866	Jan 06, 2036	U-4047		
		8580841	Mar 05, 2030	DS DP		
		9126932	Jul 22, 2029	U-3732		
<u>EVEROLIMUS - AFINITOR</u>						
N 022334	001	8410131	Nov 01, 2025	U-1368		
		8410131*PED	May 01, 2026			
		9006224	Jul 01, 2028	U-1681		
<u>EVEROLIMUS - AFINITOR</u>						
N 022334	002	8410131	Nov 01, 2025	U-1368		
		8410131*PED	May 01, 2026			
		9006224	Jul 01, 2028	U-1681		
<u>EVEROLIMUS - AFINITOR</u>						
N 022334	003	8410131	Nov 01, 2025	U-1368		
		8410131*PED	May 01, 2026			
		9006224	Jul 01, 2028	U-1681		
<u>EVEROLIMUS - AFINITOR</u>						
N 022334	004	8410131	Nov 01, 2025	U-1368		
		8410131*PED	May 01, 2026			
		9006224	Jul 01, 2028	U-1681		
<u>EVEROLIMUS - AFINITOR DISPERZ</u>						
N 203985	001				ODE-169	Apr 10, 2025
<u>EVEROLIMUS - AFINITOR DISPERZ</u>						
N 203985	002				ODE-169	Apr 10, 2025
<u>EVEROLIMUS - AFINITOR DISPERZ</u>						
N 203985	003				ODE-169	Apr 10, 2025
<u>EXENATIDE SYNTHETIC - BYDUREON</u>						
N 022200	001	6515117	Oct 04, 2025	DS DP U-2588	NPP	Jul 22, 2024
		6515117*PED	Apr 04, 2026		PED	Jan 22, 2025

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>EXENATIDE SYNTHETIC - BYDUREON</u>						
N 022200 001	7456254	Jun 30, 2025	DP U-2588			
	7456254	Jun 30, 2025	DP U-2589			
	7456254	Jun 30, 2025	DP U-2590			
	7456254	Jun 30, 2025	DP U-3188			
	7456254	Jun 30, 2025	DP U-3189			
	7456254	Jun 30, 2025	DP U-3190			
	7456254*PED	Dec 30, 2025				
	7612176	Apr 13, 2025	DP U-2588			
	7612176	Apr 13, 2025	DP U-2589			
	7612176	Apr 13, 2025	DP U-2590			
	7612176	Apr 13, 2025	DP U-3188			
	7612176	Apr 13, 2025	DP U-3189			
	7612176	Apr 13, 2025	DP U-3190			
	7612176*PED	Oct 13, 2025				
	8329648	Aug 18, 2026	U-2588			
	8329648	Aug 18, 2026	U-2589			
	8329648	Aug 18, 2026	U-2590			
	8329648	Aug 18, 2026	U-2593			
	8329648	Aug 18, 2026	U-2594			
	8329648	Aug 18, 2026	U-2595			
	8329648	Aug 18, 2026	U-2596			
	8329648	Aug 18, 2026	U-3188			
	8329648	Aug 18, 2026	U-3189			
	8329648	Aug 18, 2026	U-3190			
	8329648*PED	Feb 18, 2027				
	8361972	Mar 21, 2028	U-2588			
	8361972*PED	Sep 21, 2028				
	8431685	Apr 13, 2025	DP U-2588			
	8431685	Apr 13, 2025	DP U-2589			
	8431685	Apr 13, 2025	DP U-2590			
	8431685	Apr 13, 2025	DP U-3188			
	8431685	Apr 13, 2025	DP U-3189			
	8431685	Apr 13, 2025	DP U-3190			
	8431685*PED	Oct 13, 2025				
	8461105	Apr 13, 2025	DP U-2588			
	8461105	Apr 13, 2025	DP U-2589			
	8461105	Apr 13, 2025	DP U-2590			
	8461105	Apr 13, 2025	DP U-3188			
	8461105	Apr 13, 2025	DP U-3189			
	8461105	Apr 13, 2025	DP U-3190			
	8461105*PED	Oct 13, 2025				
	8501698	Jun 20, 2027	DP U-2588			
	8501698*PED	Dec 20, 2027				
	8906851	Aug 18, 2026	U-2588			
	8906851	Aug 18, 2026	U-2589			
	8906851	Aug 18, 2026	U-2590			
	8906851	Aug 18, 2026	U-2593			
	8906851	Aug 18, 2026	U-3188			
	8906851	Aug 18, 2026	U-3189			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>EXENATIDE SYNTHETIC - BYDUREON</u>						
N 022200 001	8906851	Aug 18, 2026	U-3190			
	8906851*PED	Feb 18, 2027				
	9884092	Aug 18, 2026	U-2588			
	9884092	Aug 18, 2026	U-2589			
	9884092	Aug 18, 2026	U-2590			
	9884092	Aug 18, 2026	U-2593			
	9884092	Aug 18, 2026	U-2594			
	9884092	Aug 18, 2026	U-2595			
	9884092	Aug 18, 2026	U-2596			
	9884092	Aug 18, 2026	U-3188			
	9884092	Aug 18, 2026	U-3189			
	9884092	Aug 18, 2026	U-3190			
	9884092*PED	Feb 18, 2027				
<u>EXENATIDE SYNTHETIC - BYDUREON PEN</u>						
N 022200 002	6515117	Oct 04, 2025	DS DP U-2588		NPP	Jul 22, 2024
	6515117*PED	Apr 04, 2026			PED	Jan 22, 2025
	7456254	Jun 30, 2025	DP U-2588			
	7456254	Jun 30, 2025	DP U-2589			
	7456254	Jun 30, 2025	DP U-2590			
	7456254	Jun 30, 2025	DP U-3188			
	7456254	Jun 30, 2025	DP U-3189			
	7456254	Jun 30, 2025	DP U-3190			
	7456254*PED	Dec 30, 2025				
	7612176	Apr 13, 2025	DP U-2588			
	7612176	Apr 13, 2025	DP U-2589			
	7612176	Apr 13, 2025	DP U-2590			
	7612176	Apr 13, 2025	DP U-3188			
	7612176	Apr 13, 2025	DP U-3189			
	7612176	Apr 13, 2025	DP U-3190			
	7612176*PED	Oct 13, 2025				
	8216180	Jan 12, 2028	DP			
	8216180*PED	Jul 12, 2028				
	8329648	Aug 18, 2026	U-2588			
	8329648	Aug 18, 2026	U-2589			
	8329648	Aug 18, 2026	U-2590			
	8329648	Aug 18, 2026	U-2593			
	8329648	Aug 18, 2026	U-2594			
	8329648	Aug 18, 2026	U-2595			
	8329648	Aug 18, 2026	U-2596			
	8329648	Aug 18, 2026	U-3188			
	8329648	Aug 18, 2026	U-3189			
	8329648	Aug 18, 2026	U-3190			
	8329648*PED	Feb 18, 2027				
	8361972	Mar 21, 2028	U-2588			
	8361972*PED	Sep 21, 2028				
	8431685	Apr 13, 2025	DP U-2588			
	8431685	Apr 13, 2025	DP U-2589			
	8431685	Apr 13, 2025	DP U-2590			
	8431685	Apr 13, 2025	DP U-3188			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>EXENATIDE SYNTHETIC - BYDUREON PEN</u>						
N 022200 002	8431685	Apr 13, 2025	DP U-3189			
	8431685	Apr 13, 2025	DP U-3190			
	8431685*PED	Oct 13, 2025				
	8439864	Mar 25, 2028	DP			
	8439864*PED	Sep 25, 2028				
	8461105	Apr 13, 2025	DP U-2588			
	8461105	Apr 13, 2025	DP U-2589			
	8461105	Apr 13, 2025	DP U-2590			
	8461105	Apr 13, 2025	DP U-3188			
	8461105	Apr 13, 2025	DP U-3189			
	8461105	Apr 13, 2025	DP U-3190			
	8461105*PED	Oct 13, 2025				
	8501698	Jun 20, 2027	DP U-2588			
	8501698*PED	Dec 20, 2027				
	8690837	May 19, 2029	DP			
	8690837*PED	Nov 19, 2029				
	8721615	Jan 18, 2030	DP			
	8721615*PED	Jul 18, 2030				
	8758292	Nov 12, 2027	DP			
	8758292*PED	May 12, 2028				
	8827963	Feb 04, 2029	DP			
	8827963*PED	Aug 04, 2029				
	8906851	Aug 18, 2026	U-2588			
	8906851	Aug 18, 2026	U-2589			
	8906851	Aug 18, 2026	U-2590			
	8906851	Aug 18, 2026	U-2593			
	8906851	Aug 18, 2026	U-3188			
	8906851	Aug 18, 2026	U-3189			
	8906851	Aug 18, 2026	U-3190			
	8906851*PED	Feb 18, 2027				
	8998876	Jan 07, 2030	DP			
	8998876*PED	Jul 07, 2030				
	9320853	Mar 25, 2028	DP			
	9320853*PED	Sep 25, 2028				
	9884092	Aug 18, 2026	U-2588			
	9884092	Aug 18, 2026	U-2589			
	9884092	Aug 18, 2026	U-2590			
	9884092	Aug 18, 2026	U-2593			
	9884092	Aug 18, 2026	U-2594			
	9884092	Aug 18, 2026	U-2595			
	9884092	Aug 18, 2026	U-2596			
	9884092	Aug 18, 2026	U-3188			
	9884092	Aug 18, 2026	U-3189			
	9884092	Aug 18, 2026	U-3190			
	9884092*PED	Feb 18, 2027				
<u>EXENATIDE SYNTHETIC - BYDUREON BCISE</u>						
N 209210 001	6515117	Oct 04, 2025	DS DP U-2588		NPP	Jul 22, 2024
	6515117*PED	Apr 04, 2026			PED	Jan 22, 2025
	7456254	Jun 30, 2025	DP U-2588			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>EXENATIDE SYNTHETIC - BYDUREON BCISE</u>						
N 209210 001	7456254	Jun 30, 2025	DP U-2589			
	7456254	Jun 30, 2025	DP U-2590			
	7456254	Jun 30, 2025	DP U-3188			
	7456254	Jun 30, 2025	DP U-3189			
	7456254	Jun 30, 2025	DP U-3190			
	7456254*PED	Dec 30, 2025				
	7612176	Apr 13, 2025	DP U-2589			
	7612176	Apr 13, 2025	DP U-2590			
	7612176	Apr 13, 2025	DP U-3188			
	7612176	Apr 13, 2025	DP U-3189			
	7612176	Apr 13, 2025	DP U-3190			
	7612176*PED	Oct 13, 2025				
	8329648	Aug 18, 2026	U-2588			
	8329648	Aug 18, 2026	U-2589			
	8329648	Aug 18, 2026	U-2590			
	8329648	Aug 18, 2026	U-2593			
	8329648	Aug 18, 2026	U-2594			
	8329648	Aug 18, 2026	U-2595			
	8329648	Aug 18, 2026	U-2596			
	8329648	Aug 18, 2026	U-3188			
	8329648	Aug 18, 2026	U-3189			
	8329648	Aug 18, 2026	U-3190			
	8329648*PED	Feb 18, 2027				
	8361972	Mar 21, 2028	U-2588			
	8361972*PED	Sep 21, 2028				
	8431685	Apr 13, 2025	DP U-2588			
	8431685	Apr 13, 2025	DP U-2589			
	8431685	Apr 13, 2025	DP U-2590			
	8431685	Apr 13, 2025	DP U-2597			
	8431685	Apr 13, 2025	DP U-3188			
	8431685	Apr 13, 2025	DP U-3189			
	8431685	Apr 13, 2025	DP U-3190			
	8431685*PED	Oct 13, 2025				
	8461105	Apr 13, 2025	DP U-2588			
	8461105	Apr 13, 2025	DP U-2589			
	8461105	Apr 13, 2025	DP U-2590			
	8461105	Apr 13, 2025	DP U-2597			
	8461105	Apr 13, 2025	DP U-3188			
	8461105	Apr 13, 2025	DP U-3189			
	8461105	Apr 13, 2025	DP U-3190			
	8461105*PED	Oct 13, 2025				
	8501698	Jun 20, 2027	U-2588			
	8501698*PED	Dec 20, 2027				
	8895033	Oct 04, 2030	DP U-2589			
	8895033	Oct 04, 2030	DP U-2590			
	8895033	Oct 04, 2030	DP U-2597			
	8895033	Oct 04, 2030	DP U-2601			
	8895033	Oct 04, 2030	DP U-2602			
	8895033	Oct 04, 2030	DP U-3188			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>EXENATIDE SYNTHETIC - BYDUREON BCISE</u>						
N 209210	001	8895033	Oct 04, 2030	DP U-3189		
		8895033	Oct 04, 2030	DP U-3190		
		8895033*PED	Apr 04, 2031			
		8906851	Aug 18, 2026	U-2588		
		8906851	Aug 18, 2026	U-2589		
		8906851	Aug 18, 2026	U-2590		
		8906851	Aug 18, 2026	U-2593		
		8906851	Aug 18, 2026	U-2597		
		8906851	Aug 18, 2026	U-3188		
		8906851	Aug 18, 2026	U-3189		
		8906851	Aug 18, 2026	U-3190		
		8906851*PED	Feb 18, 2027			
		9884092	Aug 18, 2026	U-2588		
		9884092	Aug 18, 2026	U-2589		
		9884092	Aug 18, 2026	U-2590		
		9884092	Aug 18, 2026	U-2593		
		9884092	Aug 18, 2026	U-2594		
		9884092	Aug 18, 2026	U-2595		
		9884092	Aug 18, 2026	U-2596		
		9884092	Aug 18, 2026	U-2597		
		9884092	Aug 18, 2026	U-3188		
		9884092	Aug 18, 2026	U-3189		
		9884092	Aug 18, 2026	U-3190		
		9884092*PED	Feb 18, 2027			
<u>EZETIMIBE - ZETIA</u>						
N 021445	001	7612058	Oct 30, 2025	U-1027		
		7612058	Oct 30, 2025	U-1173		
		7612058*PED	Apr 30, 2026			
<u>EZETIMIBE; ROSUVASTATIN CALCIUM - ROSZET</u>						
N 213072	001	10376470	May 01, 2033	DP U-3095		
		9763885	May 01, 2033	DP U-3095		
<u>EZETIMIBE; ROSUVASTATIN CALCIUM - ROSZET</u>						
N 213072	002	10376470	May 01, 2033	DP U-3095		
		9763885	May 01, 2033	DP U-3095		
<u>EZETIMIBE; ROSUVASTATIN CALCIUM - ROSZET</u>						
N 213072	003	10376470	May 01, 2033	DP U-3095		
		9763885	May 01, 2033	DP U-3095		
<u>EZETIMIBE; ROSUVASTATIN CALCIUM - ROSZET</u>						
N 213072	004	10376470	May 01, 2033	DP U-3095		
		9763885	May 01, 2033	DP U-3095		
<u>FAMOTIDINE; IBUPROFEN - DUEXIS</u>						
N 022519	001	8067451	Jul 18, 2026	DP U-1196		
		8309127	Jul 18, 2026	DP		
		8318202	Jul 18, 2026	DP		
		8449910	Jul 18, 2026	DP		
		8501228	Jul 18, 2026	U-1196		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>FEBUXOSTAT - ULORIC</u>						
N 021856	001 8372872	Sep 08, 2031	U-1346			
	9107912	Sep 08, 2031	U-1346			
<u>FEBUXOSTAT - ULORIC</u>						
N 021856	002 8372872	Sep 08, 2031	U-1346			
	9107912	Sep 08, 2031	U-1346			
<u>FEDRATINIB HYDROCHLORIDE - INREBIC</u>						
N 212327	001 10391094	Jun 04, 2032	DP U-2607		ODE-259	Aug 16, 2026
	11400092	Sep 24, 2039	U-3409			
	7528143	Nov 16, 2031	DS DP			
	7825246	Dec 16, 2026	DS			
	8138199	Jun 30, 2028	U-2607			
<u>FENFLURAMINE HYDROCHLORIDE - FINTEPLA</u>						
N 212102	001 10452815	Jun 29, 2038	U-2859	Y	I-887	Mar 25, 2025
	10452815*PED	Dec 29, 2038			ODE-312	Jun 25, 2027
	10478441	May 03, 2033	U-2860		ODE-393	Mar 25, 2029
	10478441*PED	Nov 03, 2033			PED	Sep 25, 2025
	10478442	May 03, 2033	U-2860		PED	Dec 25, 2027
	10478442*PED	Nov 03, 2033			PED	Sep 25, 2029
	10603290	Aug 02, 2037	U-2861			
	10603290	Aug 02, 2037	U-3347			
	10603290*PED	Feb 02, 2038				
	10947183	Dec 20, 2036	DS DP			
	10947183*PED	Jun 20, 2037				
	11040018	Aug 02, 2037	U-2861			
	11040018	Aug 02, 2037	U-3347			
	11040018*PED	Feb 02, 2038				
	11406606	Aug 02, 2037	U-3406			
	11406606	Aug 02, 2037	U-3407			
	11406606*PED	Feb 02, 2038				
	11759440	Aug 02, 2037	U-3694			
	11759440*PED	Feb 02, 2038				
	11786487	Aug 02, 2037	U-3733			
	11786487*PED	Feb 02, 2038				
	12097206	May 03, 2033	U-4013			
	12097206*PED	Nov 03, 2033				
	9549909	May 03, 2033	U-2858			
	9549909*PED	Nov 03, 2033				
	9603814	May 03, 2033	U-2858			
	9603814*PED	Nov 03, 2033				
	9603815	May 03, 2033	U-2858			
	9603815*PED	Nov 03, 2033				
	9610260	May 03, 2033	U-2858			
	9610260*PED	Nov 03, 2033				
<u>FENOFIBRATE - ANTARA (MICRONIZED)</u>						
N 021695	004 8026281	Apr 22, 2025	U-1447			
	8026281	Apr 22, 2025	U-1448			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>FENOFIBRATE - ANTARA (MICRONIZED)</u>						
N 021695	005	8026281	Apr 22, 2025			U-1447
		8026281	Apr 22, 2025			U-1448
		9314447	May 31, 2033	DP		U-1447
		9314447	May 31, 2033	DP		U-1448
<u>FENOFIBRIC ACID - FIBRICOR</u>						
N 022418	001	7569612	Aug 20, 2027			U-1000
		7741373	Aug 20, 2027			U-1059
		7741374	Aug 20, 2027			U-1060
		7741374	Aug 20, 2027			U-1061
		7915247	Aug 20, 2027			U-1000
		7915247	Aug 20, 2027			U-1059
		7915247	Aug 20, 2027			U-1061
<u>FENOFIBRIC ACID - FIBRICOR</u>						
N 022418	002	7569612	Aug 20, 2027			U-1000
		7741373	Aug 20, 2027			U-1059
		7741374	Aug 20, 2027			U-1060
		7741374	Aug 20, 2027			U-1061
		7915247	Aug 20, 2027			U-1000
		7915247	Aug 20, 2027			U-1059
		7915247	Aug 20, 2027			U-1061
<u>FENTANYL - SUBSYS</u>						
N 202788	001	10016403	Jan 25, 2027	DP		
		10610523	Jan 25, 2027	DP		
		8486972	Apr 27, 2030	DP		
		8486973	Apr 27, 2030			U-55
		8835459	Jan 25, 2027	DP		
		8835460	Jan 25, 2027	DP		U-55
		9241935	Jan 25, 2027	DP		
		9289387	Jan 25, 2027	DP		U-55
		9642797	Jan 25, 2027	DP		U-55
		9642844	Jan 25, 2027	DP		
<u>FENTANYL - SUBSYS</u>						
N 202788	002	10016403	Jan 25, 2027	DP		
		10610523	Jan 25, 2027	DP		
		8486972	Apr 27, 2030	DP		
		8486973	Apr 27, 2030			U-55
		8835460	Jan 25, 2027	DP		U-55
		9241935	Jan 25, 2027	DP		
		9289387	Jan 25, 2027	DP		U-55
		9642797	Jan 25, 2027	DP		U-55
		9642844	Jan 25, 2027	DP		
<u>FENTANYL - SUBSYS</u>						
N 202788	003	10016403	Jan 25, 2027	DP		
		10610523	Jan 25, 2027	DP		
		8486972	Apr 27, 2030	DP		
		8486973	Apr 27, 2030			U-55
		8835459	Jan 25, 2027	DP		
		8835460	Jan 25, 2027	DP		U-55

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>FENTANYL - SUBSYS</u>						
N 202788 003	9241935	Jan 25, 2027	DP			
	9289387	Jan 25, 2027	DP U-55			
	9642797	Jan 25, 2027	DP U-55			
	9642844	Jan 25, 2027	DP			
<u>FENTANYL - SUBSYS</u>						
N 202788 004	10016403	Jan 25, 2027	DP			
	10610523	Jan 25, 2027	DP			
	8486972	Apr 27, 2030	DP			
	8486973	Apr 27, 2030	U-55			
	8835459	Jan 25, 2027	DP			
	8835460	Jan 25, 2027	DP U-55			
	9241935	Jan 25, 2027	DP			
	9289387	Jan 25, 2027	DP U-55			
	9642797	Jan 25, 2027	DP U-55			
	9642844	Jan 25, 2027	DP			
<u>FENTANYL - SUBSYS</u>						
N 202788 005	10016403	Jan 25, 2027	DP			
	10610523	Jan 25, 2027	DP			
	8486972	Apr 27, 2030	DP			
	8486973	Apr 27, 2030	U-55			
	8835460	Jan 25, 2027	DP U-55			
	9241935	Jan 25, 2027	DP			
	9289387	Jan 25, 2027	DP U-55			
	9642797	Jan 25, 2027	DP U-55			
	9642844	Jan 25, 2027	DP			
<u>FENTANYL - SUBSYS</u>						
N 202788 006	10016403	Jan 25, 2027	DP			
	10610523	Jan 25, 2027	DP			
	8486972	Apr 27, 2030	DP			
	8486973	Apr 27, 2030	U-55			
	8835459	Jan 25, 2027	DP			
	8835460	Jan 25, 2027	DP U-55			
	9241935	Jan 25, 2027	DP			
	9289387	Jan 25, 2027	DP U-55			
	9642797	Jan 25, 2027	DP U-55			
	9642844	Jan 25, 2027	DP			
<u>FENTANYL - SUBSYS</u>						
N 202788 007	10016403	Jan 25, 2027	DP			
	10610523	Jan 25, 2027	DP			
	8486972	Apr 27, 2030	DP			
	8486973	Apr 27, 2030	U-55			
	8835459	Jan 25, 2027	DP			
	8835460	Jan 25, 2027	DP U-55			
	9241935	Jan 25, 2027	DP			
	9289387	Jan 25, 2027	DP U-55			
	9642797	Jan 25, 2027	DP U-55			
	9642844	Jan 25, 2027	DP			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>FENTANYL CITRATE - FENTORA</u>						
N 021947	001 7862832	Jun 15, 2028	DP			
	7862833	Jun 15, 2028	DP			
<u>FENTANYL CITRATE - FENTORA</u>						
N 021947	002 7862832	Jun 15, 2028	DP			
	7862833	Jun 15, 2028	DP			
<u>FENTANYL CITRATE - FENTORA</u>						
N 021947	003 7862832	Jun 15, 2028	DP			
	7862833	Jun 15, 2028	DP			
<u>FENTANYL CITRATE - FENTORA</u>						
N 021947	004 7862832	Jun 15, 2028	DP			
	7862833	Jun 15, 2028	DP			
<u>FENTANYL CITRATE - FENTORA</u>						
N 021947	005 7862832	Jun 15, 2028	DP			
	7862833	Jun 15, 2028	DP			
<u>FENTANYL CITRATE - ONSOLIS</u>						
N 022266	001 9597288	Jul 23, 2027	DP U-767			
<u>FENTANYL CITRATE - ONSOLIS</u>						
N 022266	002 9597288	Jul 23, 2027	DP U-767			
<u>FENTANYL CITRATE - ONSOLIS</u>						
N 022266	003 9597288	Jul 23, 2027	DP U-767			
<u>FENTANYL CITRATE - ONSOLIS</u>						
N 022266	004 9597288	Jul 23, 2027	DP U-767			
<u>FENTANYL CITRATE - ONSOLIS</u>						
N 022266	005 9597288	Jul 23, 2027	DP U-767			
<u>FENTANYL CITRATE - LAZANDA</u>						
N 022569	001 9731869	Jan 26, 2032	DP			
<u>FENTANYL CITRATE - LAZANDA</u>						
N 022569	002 9731869	Jan 26, 2032	DP			
<u>FENTANYL CITRATE - LAZANDA</u>						
N 022569	003 9731869	Jan 26, 2032	DP			
<u>FENTANYL HYDROCHLORIDE - IONSYS</u>						
N 021338	001 8301238	Sep 30, 2031	DP			
	8428708	May 21, 2032	U-736			
	8428709	Jun 11, 2032	DP U-736			
	8781571	Mar 31, 2032	DP U-736			
	9095706	Feb 03, 2033	DP			
	9364656	Sep 30, 2031	U-736			
	9731121	Oct 17, 2031	DP			
<u>FERRIC CARBOXYMALTOSE - INJECTAFER</u>						
N 203565	001 11364260	Jan 08, 2027	U-3637		I-915	May 31, 2026
	11433091	Jan 08, 2027	U-3435			
	11433091	Jan 08, 2027	U-3436			
	11433091	Jan 08, 2027	U-3437			
	11433091	Jan 08, 2027	U-3438			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>FERRIC CARBOXYMALTOSE - INJECTAFER</u>						
N 203565 001	11433091	Jan 08, 2027	U-3634			
	11478502	Jan 08, 2027	U-3472			
	11478502	Jan 08, 2027	U-3473			
	11478502	Jan 08, 2027	U-3474			
	7612109	Feb 05, 2025	DS DP			
	7754702	Feb 15, 2028	U-1432			
	7754702	Feb 15, 2028	U-3312			
	7754702	Feb 15, 2028	U-3313			
	7754702	Feb 15, 2028	U-3314			
	7754702	Feb 15, 2028	U-3636			
	8895612	Jan 08, 2027	U-1620			
	8895612	Jan 08, 2027	U-3115			
	8895612	Jan 08, 2027	U-3116			
	8895612	Jan 08, 2027	U-3315			
	8895612	Jan 08, 2027	U-3316			
	8895612	Jan 08, 2027	U-3635			
<u>FERRIC CARBOXYMALTOSE - INJECTAFER</u>						
N 203565 002	11364260	Jan 08, 2027	U-3637		I-915	May 31, 2026
	11433091	Jan 08, 2027	U-3435			
	11433091	Jan 08, 2027	U-3436			
	11433091	Jan 08, 2027	U-3437			
	11433091	Jan 08, 2027	U-3438			
	11433091	Jan 08, 2027	U-3634			
	11478502	Jan 08, 2027	U-3472			
	11478502	Jan 08, 2027	U-3473			
	11478502	Jan 08, 2027	U-3474			
	7612109	Feb 05, 2025	DS DP			
	7754702	Feb 15, 2028	U-2555			
	7754702	Feb 15, 2028	U-2556			
	7754702	Feb 15, 2028	U-2557			
	7754702	Feb 15, 2028	U-3312			
	7754702	Feb 15, 2028	U-3313			
	7754702	Feb 15, 2028	U-3314			
	7754702	Feb 15, 2028	U-3636			
	8895612	Jan 08, 2027	U-1620			
	8895612	Jan 08, 2027	U-3050			
	8895612	Jan 08, 2027	U-3051			
	8895612	Jan 08, 2027	U-3115			
	8895612	Jan 08, 2027	U-3116			
	8895612	Jan 08, 2027	U-3315			
	8895612	Jan 08, 2027	U-3316			
	8895612	Jan 08, 2027	U-3635			
<u>FERRIC CARBOXYMALTOSE - INJECTAFER</u>						
N 203565 003	11364260	Jan 08, 2027	U-3637		I-915	May 31, 2026
	11433091	Jan 08, 2027	U-3435			
	11433091	Jan 08, 2027	U-3436			
	11433091	Jan 08, 2027	U-3437			
	11433091	Jan 08, 2027	U-3438			
	11433091	Jan 08, 2027	U-3634			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>FERRIC CARBOXYMALTOSE - INJECTAFER</u>						
N 203565 003	11478502	Jan 08, 2027	U-3472			
	11478502	Jan 08, 2027	U-3473			
	11478502	Jan 08, 2027	U-3474			
	7612109	Feb 05, 2025	DS DP			
	7754702	Feb 15, 2028	U-2555			
	7754702	Feb 15, 2028	U-2556			
	7754702	Feb 15, 2028	U-2557			
	7754702	Feb 15, 2028	U-3312			
	7754702	Feb 15, 2028	U-3313			
	7754702	Feb 15, 2028	U-3314			
	7754702	Feb 15, 2028	U-3636			
	8895612	Jan 08, 2027	U-1620			
	8895612	Jan 08, 2027	U-3050			
	8895612	Jan 08, 2027	U-3051			
	8895612	Jan 08, 2027	U-3115			
	8895612	Jan 08, 2027	U-3116			
	8895612	Jan 08, 2027	U-3315			
	8895612	Jan 08, 2027	U-3316			
	8895612	Jan 08, 2027	U-3635			
<u>FERRIC CARBOXYMALTOSE - INJECTAFER</u>						
N 203565 004	11364260	Jan 08, 2027	U-3637		I-915	May 31, 2026
	11433091	Jan 08, 2027	U-3435			
	11433091	Jan 08, 2027	U-3436			
	11433091	Jan 08, 2027	U-3437			
	11433091	Jan 08, 2027	U-3438			
	11433091	Jan 08, 2027	U-3634			
	11478502	Jan 08, 2027	U-3472			
	11478502	Jan 08, 2027	U-3473			
	11478502	Jan 08, 2027	U-3474			
	7612109	Feb 05, 2025	DS DP			
	7754702	Feb 15, 2028	U-3312			
	7754702	Feb 15, 2028	U-3313			
	7754702	Feb 15, 2028	U-3314			
	7754702	Feb 15, 2028	U-3636			
	8895612	Jan 08, 2027	U-3115			
	8895612	Jan 08, 2027	U-3116			
	8895612	Jan 08, 2027	U-3315			
	8895612	Jan 08, 2027	U-3316			
	8895612	Jan 08, 2027	U-3635			
<u>FERRIC CITRATE - AURYXIA</u>						
N 205874 001	10300039	Jul 21, 2030	U-2549			
	8093423	Apr 21, 2026	U-1577			
	9387191	Jul 21, 2030	DP			
<u>FERRIC DERISOMALTOSE - MONOFERRIC</u>						
N 208171 001	10414831	Mar 25, 2029	DS DP			
	11633489	Jun 22, 2036	U-3594			
	11851504	Mar 25, 2029	DS DP			
	12030962	Mar 25, 2029	DS DP			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>FERRIC DERISOMALTOSE - MONOFERRIC</u>						
N 208171	001 8815301	Aug 14, 2029	DS DP U-2734			
<u>FERRIC DERISOMALTOSE - MONOFERRIC</u>						
N 208171	002 10414831	Mar 25, 2029	DS DP			
	11633489	Jun 22, 2036			U-3594	
	11851504	Mar 25, 2029	DS DP			
	12030962	Mar 25, 2029	DS DP			
	8815301	Aug 14, 2029	DS DP U-2734			
<u>FERRIC DERISOMALTOSE - MONOFERRIC</u>						
N 208171	003 10414831	Mar 25, 2029	DS DP			
	11633489	Jun 22, 2036			U-3594	
	11851504	Mar 25, 2029	DS DP			
	12030962	Mar 25, 2029	DS DP			
	8815301	Aug 14, 2029	DS DP U-2734			
<u>FERRIC MALTOL - ACCRUFER</u>						
N 212320	001 10179120	Jan 06, 2035			U-2603	
	9248148	Mar 29, 2031			U-2603	
	9802973	Oct 23, 2035	DS DP U-2603			
<u>FERRIC OXYHYDROXIDE - VELPHORO</u>						
N 205109	001 10624855	Nov 26, 2034	DP		NPP	Jul 01, 2027
	10624855*PED	May 26, 2035			PED	Jan 01, 2028
	10682376	Nov 13, 2028	DP			
	10682376*PED	May 13, 2029				
	10695367	Nov 13, 2028	DP			
	10695367*PED	May 13, 2029				
	10925896	Nov 13, 2028	DP			
	10925896*PED	May 13, 2029				
	10925897	Nov 13, 2028	DP			
	10925897*PED	May 13, 2029				
	10933090	Nov 13, 2028	DP			
	10933090*PED	May 13, 2029				
	11013761	Nov 13, 2028	DP			
	11013761*PED	May 13, 2029				
	11013762	Nov 13, 2028	DP			
	11013762*PED	May 13, 2029				
	11234938	Nov 26, 2034	DP			
	11234938*PED	May 26, 2035				
	11446252	Nov 26, 2034	DP			
	11446252*PED	May 26, 2035				
	9561251	Jan 23, 2030	DP U-1468			
	9561251*PED	Jul 23, 2030				
<u>FERRIC PYROPHOSPHATE CITRATE - TRIFERIC</u>						
N 206317	001 7816404	Apr 17, 2029	DP U-1656			
<u>FERRIC PYROPHOSPHATE CITRATE - TRIFERIC</u>						
N 208551	001 7816404	Apr 17, 2029	U-1656			
	7857977	Sep 08, 2027	U-1656			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>FERRIC PYROPHOSPHATE CITRATE - TRIFERIC AVNU</u>						
N 212860	001 7816404	Apr 17, 2029	DS U-2801			
<u>FESOTERODINE FUMARATE - TOVIAZ</u>						
N 022030	001 7807715	Jun 07, 2027	DP U-913			
	8088398	Jun 07, 2027	DP U-913			
	8501723	Jun 07, 2027	DP			
	8501723*PED	Dec 07, 2027				
<u>FESOTERODINE FUMARATE - TOVIAZ</u>						
N 022030	002 7807715	Jun 07, 2027	DP U-913			
	8088398	Jun 07, 2027	DP U-913			
	8501723	Jun 07, 2027	DP			
	8501723*PED	Dec 07, 2027				
<u>FEXINIDAZOLE - FEXINIDAZOLE</u>						
N 214429	001				NCE	Jul 16, 2026
					ODE-359	Jul 16, 2028
<u>FEXOFENADINE HYDROCHLORIDE - CHILDREN'S ALLEGRA ALLERGY</u>						
N 201373	001 8933097	Aug 02, 2030	DP			
<u>FEXOFENADINE HYDROCHLORIDE - CHILDREN'S ALLEGRA HIVES</u>						
N 201373	002 8933097	Aug 02, 2030	DP			
<u>FEZOLINETANT - VEOZAH</u>						
N 216578	001 10836768	Mar 28, 2034	U-3621		NCE	May 12, 2028
	8871761	Apr 04, 2031	DS			
	9422299	Mar 28, 2034	DS DP U-3622			
	9987274	Mar 28, 2034	U-3622			
<u>FIDAXOMICIN - DIFICID</u>						
N 201699	001 7378508	Jul 31, 2027	DS DP		ODE-367	Jan 24, 2027
	7863249	Jul 31, 2027	DS DP			
	7906489	Mar 04, 2027	U-2741			
	7906489	Mar 04, 2027	U-319			
	7906489*PED	Sep 04, 2027				
	8859510	Jul 31, 2027	U-2741			
	8859510	Jul 31, 2027	U-319			
	8859510*PED	Jan 31, 2028				
<u>FIDAXOMICIN - DIFICID</u>						
N 213138	001 7378508	Jul 31, 2027	DS DP		ODE-367	Jan 24, 2027
	7378508*PED	Jan 31, 2028				
	7863249	Jul 31, 2027	DP			
	7863249*PED	Jan 31, 2028				
	7906489	Mar 04, 2027	U-2741			
	7906489*PED	Sep 04, 2027				
	8859510	Jul 31, 2027	U-2741			
	8859510*PED	Jan 31, 2028				
	9808530	May 28, 2034	DP			
	9808530*PED	Nov 28, 2034				
<u>FINAFLOXACIN - XTORO</u>						
N 206307	001 8536167	Aug 08, 2031	U-1679			
	9119859	Jul 02, 2030	U-1679			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>FINAFLOXACIN - XTORO</u>						
N 206307	001 9504691	Nov 21, 2033	DP U-1679			
	9993483	Jul 02, 2030	DP			
<u>FINERENONE - KERENDIA</u>						
N 215341	001 8436180	Apr 12, 2029	DS DP		M-279	Sep 01, 2025
	RE49826	Jul 29, 2035	DS		NCE	Jul 09, 2026
<u>FINERENONE - KERENDIA</u>						
N 215341	002 8436180	Apr 12, 2029	DS DP		M-279	Sep 01, 2025
	RE49826	Jul 29, 2035	DS		NCE	Jul 09, 2026
<u>FINGOLIMOD HYDROCHLORIDE - GILENYA</u>						
N 022527	001 10543179	Dec 25, 2027		U-2719		
	8324283	Mar 29, 2026	DP			
	8324283*PED	Sep 29, 2026				
	9187405	Jun 25, 2027		U-2613		
	9187405*PED	Dec 25, 2027				
<u>FINGOLIMOD HYDROCHLORIDE - GILENYA</u>						
N 022527	002 9592208	Mar 30, 2032	DP U-2315			
	9592208*PED	Sep 30, 2032				
<u>FINGOLIMOD LAURYL SULFATE - TASCENSO ODT</u>						
N 214962	001 10555902	Jan 19, 2036		U-3268		
	10925829	Jan 19, 2036	DP			
	9925138	Jan 19, 2036	DP			
<u>FINGOLIMOD LAURYL SULFATE - TASCENSO ODT</u>						
N 214962	002 10555902	Jan 19, 2036		U-3493		
	10925829	Jan 19, 2036	DP			
	9925138	Jan 19, 2036	DP			
<u>FISH OIL TRIGLYCERIDES - OMEGAVEN</u>						
N 210589	001 9566260	Jul 11, 2025	DP U-2366		ODE-202	Jul 27, 2025
	9629821	Jul 11, 2025	DP U-2367			
<u>FISH OIL TRIGLYCERIDES - OMEGAVEN</u>						
N 210589	002 9566260	Jul 11, 2025	DP U-2366		ODE-202	Jul 27, 2025
	9629821	Jul 11, 2025	DP U-2367			
<u>FISH OIL; MEDIUM CHAIN TRIGLYCERIDES; OLIVE OIL; SOYBEAN OIL - SMOFLIPID 20%</u>						
N 207648	001				NPP	Mar 22, 2025
<u>FISH OIL; MEDIUM CHAIN TRIGLYCERIDES; OLIVE OIL; SOYBEAN OIL - SMOFLIPID 20%</u>						
N 207648	002				NPP	Mar 22, 2025
<u>FISH OIL; MEDIUM CHAIN TRIGLYCERIDES; OLIVE OIL; SOYBEAN OIL - SMOFLIPID 20%</u>						
N 207648	003				NPP	Mar 22, 2025
<u>FISH OIL; MEDIUM CHAIN TRIGLYCERIDES; OLIVE OIL; SOYBEAN OIL - SMOFLIPID 20%</u>						
N 207648	004				NPP	Mar 22, 2025
<u>FLIBANSERIN - ADDYI</u>						
N 022526	001 7151103	May 09, 2028	U-1734			
<u>FLORBETABEN F-18 - NEURACEQ</u>						
N 204677	001 7807135	Mar 18, 2029	DS DP U-1497			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>FLORBETAPIR F-18 - AMYVID</u>						
N 202008	001 7687052	Apr 30, 2027	DS DP			
	8506929	Apr 30, 2027	DS DP U-1423			
<u>FLORBETAPIR F-18 - AMYVID</u>						
N 202008	002 7687052	Apr 30, 2027	DS DP			
	8506929	Apr 30, 2027	DS DP U-1423			
<u>FLORBETAPIR F-18 - AMYVID</u>						
N 202008	003 7687052	Apr 30, 2027	DS DP			
	8506929	Apr 30, 2027	DS DP U-1423			
<u>FLORTAUCIPIR F-18 - TAUVID</u>						
N 212123	001 8932557	May 26, 2032	DS		NCE	May 28, 2025
<u>FLORTAUCIPIR F-18 - TAUVID</u>						
N 212123	002 8932557	May 26, 2032	DS		NCE	May 28, 2025
<u>FLORTAUCIPIR F-18 - TAUVID</u>						
N 212123	003 8932557	May 26, 2032	DS			
<u>FLORTAUCIPIR F-18 - TAUVID</u>						
N 212123	004 8932557	May 26, 2032	DS			
<u>FLOTUFOLASTAT F-18 GALLIUM - POSLUMA</u>						
N 216023	001 11413360	Nov 22, 2038	DS U-3614		NCE	May 25, 2028
	11413360	Nov 22, 2038	DS U-3615			
	12036290	Nov 27, 2041	DP U-3614			
	12036290	Nov 27, 2041	DP U-3615			
<u>FLUCICLOVINE F-18 - AXUMIN</u>						
N 208054	001 10010632	Nov 28, 2026	DP			
	10124079	Dec 30, 2035	U-2450			
	10716868	Dec 30, 2035	U-2450			
	10933147	Dec 30, 2035	U-2450			
	10953112	Nov 28, 2026	U-1879			
	10967077	Dec 30, 2035	U-2450			
	11980674	Apr 23, 2042	U-2450			
	9387266	Nov 28, 2026	U-1879			
<u>FLUOCINOLONE ACETONIDE - IUVIEN</u>						
N 201923	001 8871241	Aug 12, 2027	DP			
<u>FLUOCINOLONE ACETONIDE - YUTIQ</u>						
N 210331	001 7998108	Jan 12, 2028	DP U-3410			
	8871241	Aug 12, 2027	DP			
<u>FLUOROESTRADIOL F-18 - CERIANNA</u>						
N 212155	001				NCE	May 20, 2025
<u>FLURPIRIDAZ F-18 - FLYRCADO</u>						
N 215168	001 7344702	May 26, 2026	DS		NCE	Sep 27, 2029
	8226929	Jun 21, 2028	U-4011			
	8936777	Jun 30, 2031	U-4011			
	9161997	Feb 04, 2026	DS DP U-4011			
	9603951	May 02, 2031	U-4011			
	9687571	Nov 01, 2032	DP U-4011			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>FLUTEMETAMOL F-18 - VIZAMYL</u>						
N 203137	001	7270800	Sep 03, 2025	DS DP U-336		
		8916131	Sep 16, 2028	DP		
<u>FLUTEMETAMOL F-18 - VIZAMYL</u>						
N 203137	002	7270800	Sep 03, 2025	DS DP U-336		
		8916131	Sep 16, 2028	DP		
<u>FLUTICASONE FUROATE - FLONASE SENSIMIST ALLERGY RELIEF</u>						
N 022051	002	8062264	Apr 05, 2026	DP		
		8147461	Oct 15, 2028	DP		
		8347879	Jul 15, 2028	DP		
<u>FLUTICASONE FUROATE - ARNUITY ELLIPTA</u>						
N 205625	001	8201556	Feb 05, 2029	DP		
		8746242	Oct 11, 2030	DP		
		8746242*PED	Apr 11, 2031			
		9333310	Oct 02, 2027	DP		
		9333310*PED	Apr 02, 2028			
<u>FLUTICASONE FUROATE - ARNUITY ELLIPTA</u>						
N 205625	002	8201556	Feb 05, 2029	DP		
		8746242	Oct 11, 2030	DP		
		8746242*PED	Apr 11, 2031			
		9333310	Oct 02, 2027	DP		
		9333310*PED	Apr 02, 2028			
<u>FLUTICASONE FUROATE - ARNUITY ELLIPTA</u>						
N 205625	003	8201556	Feb 05, 2029	DP	M-290	Mar 01, 2026
		8746242	Oct 11, 2030	DP		
		8746242*PED	Apr 11, 2031			
		9333310	Oct 02, 2027	DP		
		9333310*PED	Apr 02, 2028			
<u>FLUTICASONE FUROATE; UMECLIDINIUM BROMIDE; VILANTEROL TRIFENATATE - TRELEGY ELLIPTA</u>						
N 209482	001	11090294	Nov 29, 2030	U-3202		
		7439393	May 21, 2025	DS DP U-2127		
		7439393	May 21, 2025	DS DP U-2957		
		7439393*PED	Nov 21, 2025			
		7488827	Dec 18, 2027	DS DP		
		7498440	Apr 27, 2025	DS DP		
		8161968	Feb 05, 2028	DP		
		8161968*PED	Aug 05, 2028			
		8183257	Jul 27, 2025		U-2128	
		8183257	Jul 27, 2025		U-2129	
		8309572	Apr 27, 2025		U-2129	
		8511304	Jun 14, 2027	DP U-2954		
		8511304*PED	Dec 14, 2027			
		8534281	Mar 08, 2030	DP		
		8534281*PED	Sep 08, 2030			
		8746242	Oct 11, 2030	DP		
		8746242*PED	Apr 11, 2031			
		9333310	Oct 02, 2027	DP		
		9333310*PED	Apr 02, 2028			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>FLUTICASONE FUROATE; UMECLIDINIUM BROMIDE; VILANTEROL TRIFENATATE - TRELEGY ELLIPTA</u>						
N 209482	001 9750726	Nov 29, 2030	DP			
<u>FLUTICASONE FUROATE; UMECLIDINIUM BROMIDE; VILANTEROL TRIFENATATE - TRELEGY ELLIPTA</u>						
N 209482	002 7439393	May 21, 2025	DS DP U-2957			
	7439393*PED	Nov 21, 2025				
	7488827	Dec 18, 2027	DS DP			
	7498440	Apr 27, 2025	DS DP			
	8161968	Feb 05, 2028	DP			
	8161968*PED	Aug 05, 2028				
	8183257	Jul 27, 2025		U-2129		
	8309572	Apr 27, 2025		U-2129		
	8511304	Jun 14, 2027	DP U-2954			
	8511304*PED	Dec 14, 2027				
	8534281	Mar 08, 2030	DP			
	8534281*PED	Sep 08, 2030				
	8746242	Oct 11, 2030	DP			
	8746242*PED	Apr 11, 2031				
	9333310	Oct 02, 2027	DP			
	9333310*PED	Apr 02, 2028				
	9750726	Nov 29, 2030	DP			
<u>FLUTICASONE FUROATE; VILANTEROL TRIFENATATE - BREO ELLIPTA</u>						
N 204275	001 11116721	Feb 26, 2029	DP U-1401		NPP	May 13, 2026
	11116721	Feb 26, 2029	DP U-1691		PED	Nov 13, 2026
	11116721	Feb 26, 2029	DP U-3623			
	11116721*PED	Aug 26, 2029				
	7439393	May 21, 2025	DS DP U-1401			
	7439393	May 21, 2025	DS DP U-1691			
	7439393	May 21, 2025	DS DP U-2099			
	7439393	May 21, 2025	DS DP U-2100			
	7439393	May 21, 2025	DS DP U-3623			
	7439393*PED	Nov 21, 2025				
	8161968	Feb 05, 2028	DP			
	8161968*PED	Aug 05, 2028				
	8511304	Jun 14, 2027	DP U-1424			
	8511304	Jun 14, 2027	DP U-1691			
	8511304	Jun 14, 2027	DP U-3623			
	8511304*PED	Dec 14, 2027				
	8534281	Mar 08, 2030	DP			
	8534281*PED	Sep 08, 2030				
	8746242	Oct 11, 2030	DP			
	8746242*PED	Apr 11, 2031				
	9333310	Oct 02, 2027	DP			
	9333310*PED	Apr 02, 2028				
<u>FLUTICASONE FUROATE; VILANTEROL TRIFENATATE - BREO ELLIPTA</u>						
N 204275	002 11116721	Feb 26, 2029	DP U-1691			
	11116721	Feb 26, 2029	DP U-3623			
	11116721*PED	Aug 26, 2029				
	7439393	May 21, 2025	DS DP U-1691			
	7439393	May 21, 2025	DS DP U-2099			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>FLUTICASONE FUROATE; VILANTEROL TRIFENATATE - BREO ELLIPTA</u>						
N 204275	002	7439393	May 21, 2025	DS DP U-2100		
		7439393	May 21, 2025	DS DP U-3623		
		7439393*PED	Nov 21, 2025			
		8161968	Feb 05, 2028	DP		
		8161968*PED	Aug 05, 2028			
		8511304	Jun 14, 2027	DP U-1691		
		8511304	Jun 14, 2027	DP U-3623		
		8511304*PED	Dec 14, 2027			
		8534281	Mar 08, 2030	DP		
		8534281*PED	Sep 08, 2030			
		8746242	Oct 11, 2030	DP		
		8746242*PED	Apr 11, 2031			
		9333310	Oct 02, 2027	DP		
		9333310*PED	Apr 02, 2028			
<u>FLUTICASONE FUROATE; VILANTEROL TRIFENATATE - BREO ELLIPTA</u>						
N 204275	003	11116721	Feb 26, 2029	DP U-3623	NS	May 13, 2026
		11116721*PED	Aug 26, 2029		PED	Nov 13, 2026
		7439393	May 21, 2025	DS DP U-3623		
		7439393*PED	Nov 21, 2025			
		8161968	Feb 05, 2028	DP		
		8161968*PED	Aug 05, 2028			
		8511304	Jun 14, 2027	DP U-3623		
		8511304*PED	Dec 14, 2027			
		8534281	Mar 08, 2030	DP		
		8534281*PED	Sep 08, 2030			
		8746242	Oct 11, 2030	DP		
		8746242*PED	Apr 11, 2031			
		9333310	Oct 02, 2027	DP		
		9333310*PED	Apr 02, 2028			
<u>FLUTICASONE PROPIONATE - FLOVENT HFA</u>						
N 021433	001	7500444	Feb 26, 2026	DP	Y	
		7500444*PED	Aug 26, 2026			
<u>FLUTICASONE PROPIONATE - FLOVENT HFA</u>						
N 021433	002	7500444	Feb 26, 2026	DP	Y	
		7500444*PED	Aug 26, 2026			
<u>FLUTICASONE PROPIONATE - ARMONAIR RESPICLICK</u>						
N 208798	001	10022510	May 18, 2031	DP	NPP	Jul 09, 2024
		10022510*PED	Nov 18, 2031		PED	Jan 09, 2025
		10124131	May 18, 2031	DP		
		10124131*PED	Nov 18, 2031			
		10195375	Feb 14, 2031	DP		
		10195375*PED	Aug 14, 2031			
		10561808	Jan 01, 2032	DP		
		10561808*PED	Jul 01, 2032			
		10765820	May 19, 2025	DP		
		10765820*PED	Nov 19, 2025			
		11969544	Aug 20, 2039	DP		
		11969544*PED	Feb 20, 2040			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>FLUTICASONE PROPIONATE - ARMONAIR RESPICLICK</u>						
N 208798 001	8651103	Mar 26, 2028	DP			
	8651103*PED	Sep 26, 2028				
	8714149	Feb 25, 2032	DP			
	8714149*PED	Aug 25, 2032				
	8978966	Jan 13, 2032	DP			
	8978966*PED	Jul 13, 2032				
	9216260	Jun 28, 2031	DP			
	9216260*PED	Dec 28, 2031				
	9463288	May 19, 2025	DP			
	9463288*PED	Nov 19, 2025				
	9616024*PED	Mar 01, 2025				
	9731087	May 18, 2031	DP			
	9731087*PED	Nov 18, 2031				
<u>FLUTICASONE PROPIONATE - ARMONAIR RESPICLICK</u>						
N 208798 002	10022510	May 18, 2031	DP			
	10022510*PED	Nov 18, 2031				
	10124131	May 18, 2031	DP			
	10124131*PED	Nov 18, 2031				
	10195375	Feb 14, 2031	DP			
	10195375*PED	Aug 14, 2031				
	10561808	Jan 01, 2032	DP			
	10561808*PED	Jul 01, 2032				
	10765820	May 19, 2025	DP			
	10765820*PED	Nov 19, 2025				
	11969544	Aug 20, 2039	DP			
	11969544*PED	Feb 20, 2040				
	8651103	Mar 26, 2028	DP			
	8651103*PED	Sep 26, 2028				
	8714149	Feb 25, 2032	DP			
	8714149*PED	Aug 25, 2032				
	8978966	Jan 13, 2032	DP			
	8978966*PED	Jul 13, 2032				
	9216260	Jun 28, 2031	DP			
	9216260*PED	Dec 28, 2031				
	9463288	May 19, 2025	DP			
	9463288*PED	Nov 19, 2025				
	9616024*PED	Mar 01, 2025				
	9731087	May 18, 2031	DP			
	9731087*PED	Nov 18, 2031				
<u>FLUTICASONE PROPIONATE - ARMONAIR RESPICLICK</u>						
N 208798 003	10022510	May 18, 2031	DP			
	10022510*PED	Nov 18, 2031				
	10124131	May 18, 2031	DP			
	10124131*PED	Nov 18, 2031				
	10195375	Feb 14, 2031	DP			
	10195375*PED	Aug 14, 2031				
	10561808	Jan 01, 2032	DP			
	10561808*PED	Jul 01, 2032				
	10765820	May 19, 2025	DP			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>FLUTICASONE PROPIONATE - ARMONAIR RESPICLICK</u>						
N 208798 003	10765820*PED	Nov 19, 2025				
	11969544	Aug 20, 2039	DP			
	11969544*PED	Feb 20, 2040				
	8651103	Mar 26, 2028	DP			
	8651103*PED	Sep 26, 2028				
	8714149	Feb 25, 2032	DP			
	8714149*PED	Aug 25, 2032				
	8978966	Jan 13, 2032	DP			
	8978966*PED	Jul 13, 2032				
	9216260	Jun 28, 2031	DP			
	9216260*PED	Dec 28, 2031				
	9463288	May 19, 2025	DP			
	9463288*PED	Nov 19, 2025				
	9616024*PED	Mar 01, 2025				
	9731087	May 18, 2031	DP			
	9731087*PED	Nov 18, 2031				
<u>FLUTICASONE PROPIONATE - ARMONAIR DIGIHALER</u>						
N 208798 004	10022510	May 18, 2031	DP		NPP	Jul 09, 2024
	10022510*PED	Nov 18, 2031			PED	Jan 09, 2025
	10124131	May 18, 2031	DP			
	10124131*PED	Nov 18, 2031				
	10195375	Feb 14, 2031	DP			
	10195375*PED	Aug 14, 2031				
	10561808	Jan 01, 2032	DP			
	10561808*PED	Jul 01, 2032				
	10569034	Aug 16, 2036	DP			
	10569034*PED	Feb 16, 2037				
	10765820	May 19, 2025	DP			
	10765820*PED	Nov 19, 2025				
	10918816	Dec 14, 2035	DP			
	10918816*PED	Jun 14, 2036				
	11000653	Dec 18, 2038	DP			
	11000653*PED	Jun 18, 2039				
	11173259	Jul 06, 2040	DP			
	11173259*PED	Jan 06, 2041				
	11266796	Feb 22, 2041	DP			
	11266796*PED	Aug 22, 2041				
	11344685	Sep 26, 2039	DP			
	11344685*PED	Mar 26, 2040				
	11351317	Feb 10, 2038	DP			
	11351317*PED	Aug 10, 2038				
	11357935	Sep 24, 2038	DP			
	11357935*PED	Mar 24, 2039				
	11439777	May 24, 2040	DP			
	11439777*PED	Nov 24, 2040				
	11464923	Jun 19, 2040	DP			
	11464923*PED	Dec 19, 2040				
	11969544	Aug 20, 2039	DP			
	11969544*PED	Feb 20, 2040				

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>FLUTICASONE PROPIONATE - ARMONAIR DIGIHALER</u>						
N 208798 004	8651103	Mar 26, 2028	DP			
	8651103*PED	Sep 26, 2028				
	8714149	Feb 25, 2032	DP			
	8714149*PED	Aug 25, 2032				
	8978966	Jan 13, 2032	DP			
	8978966*PED	Jul 13, 2032				
	9216260	Jun 28, 2031	DP			
	9216260*PED	Dec 28, 2031				
	9463288	May 19, 2025	DP			
	9463288*PED	Nov 19, 2025				
	9616024*PED	Mar 01, 2025				
	9731087	May 18, 2031	DP			
	9731087*PED	Nov 18, 2031				
	9782550	Aug 28, 2035	DP			
	9782550*PED	Feb 28, 2036				
	9782551	Aug 28, 2035	DP			
	9782551*PED	Feb 28, 2036				
<u>FLUTICASONE PROPIONATE - ARMONAIR DIGIHALER</u>						
N 208798 005	10022510	May 18, 2031	DP			
	10022510*PED	Nov 18, 2031				
	10124131	May 18, 2031	DP			
	10124131*PED	Nov 18, 2031				
	10195375	Feb 14, 2031	DP			
	10195375*PED	Aug 14, 2031				
	10561808	Jan 01, 2032	DP			
	10561808*PED	Jul 01, 2032				
	10569034	Aug 16, 2036	DP			
	10569034*PED	Feb 16, 2037				
	10765820	May 19, 2025	DP			
	10765820*PED	Nov 19, 2025				
	10918816	Dec 14, 2035	DP			
	10918816*PED	Jun 14, 2036				
	11000653	Dec 18, 2038	DP			
	11000653*PED	Jun 18, 2039				
	11173259	Jul 06, 2040	DP			
	11173259*PED	Jan 06, 2041				
	11266796	Feb 22, 2041	DP			
	11266796*PED	Aug 22, 2041				
	11344685	Sep 26, 2039	DP			
	11344685*PED	Mar 26, 2040				
	11351317	Feb 10, 2038	DP			
	11351317*PED	Aug 10, 2038				
	11357935	Sep 24, 2038	DP			
	11357935*PED	Mar 24, 2039				
	11439777	May 24, 2040	DP			
	11439777*PED	Nov 24, 2040				
	11464923	Jun 19, 2040	DP			
	11464923*PED	Dec 19, 2040				
	11969544	Aug 20, 2039	DP			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>FLUTICASONE PROPIONATE - ARMONAIR DIGIHALER</u>						
N 208798 005	11969544*PED	Feb 20, 2040				
	8651103	Mar 26, 2028	DP			
	8651103*PED	Sep 26, 2028				
	8714149	Feb 25, 2032	DP			
	8714149*PED	Aug 25, 2032				
	8978966	Jan 13, 2032	DP			
	8978966*PED	Jul 13, 2032				
	9216260	Jun 28, 2031	DP			
	9216260*PED	Dec 28, 2031				
	9463288	May 19, 2025	DP			
	9463288*PED	Nov 19, 2025				
	9616024*PED	Mar 01, 2025				
	9731087	May 18, 2031	DP			
	9731087*PED	Nov 18, 2031				
	9782550	Aug 28, 2035	DP			
	9782550*PED	Feb 28, 2036				
	9782551	Aug 28, 2035	DP			
	9782551*PED	Feb 28, 2036				
<u>FLUTICASONE PROPIONATE - ARMONAIR DIGIHALER</u>						
N 208798 006	10022510	May 18, 2031	DP			
	10022510*PED	Nov 18, 2031				
	10124131	May 18, 2031	DP			
	10124131*PED	Nov 18, 2031				
	10195375	Feb 14, 2031	DP			
	10195375*PED	Aug 14, 2031				
	10561808	Jan 01, 2032	DP			
	10561808*PED	Jul 01, 2032				
	10569034	Aug 16, 2036	DP			
	10569034*PED	Feb 16, 2037				
	10765820	May 19, 2025	DP			
	10765820*PED	Nov 19, 2025				
	10918816	Dec 14, 2035	DP			
	10918816*PED	Jun 14, 2036				
	11000653	Dec 18, 2038	DP			
	11000653*PED	Jun 18, 2039				
	11173259	Jul 06, 2040	DP			
	11173259*PED	Jan 06, 2041				
	11266796	Feb 22, 2041	DP			
	11266796*PED	Aug 22, 2041				
	11344685	Sep 26, 2039	DP			
	11344685*PED	Mar 26, 2040				
	11351317	Feb 10, 2038	DP			
	11351317*PED	Aug 10, 2038				
	11357935	Sep 24, 2038	DP			
	11357935*PED	Mar 24, 2039				
	11439777	May 24, 2040	DP			
	11439777*PED	Nov 24, 2040				
	11464923	Jun 19, 2040	DP			
	11464923*PED	Dec 19, 2040				

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>FLUTICASONE PROPIONATE - ARMONAIR DIGIHALER</u>						
N 208798 006	11969544	Aug 20, 2039	DP			
	11969544*PED	Feb 20, 2040				
	8651103	Mar 26, 2028	DP			
	8651103*PED	Sep 26, 2028				
	8714149	Feb 25, 2032	DP			
	8714149*PED	Aug 25, 2032				
	8978966	Jan 13, 2032	DP			
	8978966*PED	Jul 13, 2032				
	9216260	Jun 28, 2031	DP			
	9216260*PED	Dec 28, 2031				
	9463288	May 19, 2025	DP			
	9463288*PED	Nov 19, 2025				
	9616024*PED	Mar 01, 2025				
	9731087	May 18, 2031	DP			
	9731087*PED	Nov 18, 2031				
	9782550	Aug 28, 2035	DP			
	9782550*PED	Feb 28, 2036				
	9782551	Aug 28, 2035	DP			
	9782551*PED	Feb 28, 2036				
<u>FLUTICASONE PROPIONATE - ARMONAIR RESPICLICK</u>						
N 208798 007	10022510	May 18, 2031	DP		NS	Jul 09, 2024
	10022510*PED	Nov 18, 2031			PED	Jan 09, 2025
	10124131	May 18, 2031	DP			
	10124131*PED	Nov 18, 2031				
	10195375	Feb 14, 2031	DP			
	10195375*PED	Aug 14, 2031				
	10561808	Jan 01, 2032	DP			
	10561808*PED	Jul 01, 2032				
	10765820	May 19, 2025	DP			
	10765820*PED	Nov 19, 2025				
	11969544	Aug 20, 2039	DP			
	11969544*PED	Feb 20, 2040				
	8651103	Mar 26, 2028	DP			
	8651103*PED	Sep 26, 2028				
	8714149	Feb 25, 2032	DP			
	8714149*PED	Aug 25, 2032				
	8978966	Jan 13, 2032	DP			
	8978966*PED	Jul 13, 2032				
	9216260	Jun 28, 2031	DP			
	9216260*PED	Dec 28, 2031				
	9463288	May 19, 2025	DP			
	9463288*PED	Nov 19, 2025				
	9616024*PED	Mar 01, 2025				
	9731087	May 18, 2031	DP			
	9731087*PED	Nov 18, 2031				
<u>FLUTICASONE PROPIONATE - ARMONAIR DIGIHALER</u>						
N 208798 008	10022510	May 18, 2031	DP			
	10022510*PED	Nov 18, 2031				
	10124131	May 18, 2031	DP			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>FLUTICASONE PROPIONATE - ARMONAIR DIGIHALER</u>						
N 208798 008	10124131*PED	Nov 18, 2031				
	10195375	Feb 14, 2031	DP			
	10195375*PED	Aug 14, 2031				
	10561808	Jan 01, 2032	DP			
	10561808*PED	Jul 01, 2032				
	10569034	Aug 16, 2036	DP			
	10569034*PED	Feb 16, 2037				
	10765820	May 19, 2025	DP			
	10765820*PED	Nov 19, 2025				
	10918816	Dec 14, 2035	DP			
	10918816*PED	Jun 14, 2036				
	11000653	Dec 18, 2038	DP			
	11000653*PED	Jun 18, 2039				
	11173259	Jul 06, 2040	DP			
	11173259*PED	Jan 06, 2041				
	11266796	Feb 22, 2041	DP			
	11266796*PED	Aug 22, 2041				
	11344685	Sep 26, 2039	DP			
	11344685*PED	Mar 26, 2040				
	11351317	Feb 10, 2038	DP			
	11351317*PED	Aug 10, 2038				
	11357935	Sep 24, 2038	DP			
	11357935*PED	Mar 24, 2039				
	11439777	May 24, 2040	DP			
	11439777*PED	Nov 24, 2040				
	11464923	Jun 19, 2040	DP			
	11464923*PED	Dec 19, 2040				
	11969544	Aug 20, 2039	DP			
	11969544*PED	Feb 20, 2040				
	8651103	Mar 26, 2028	DP			
	8651103*PED	Sep 26, 2028				
	8714149	Nov 08, 2030	DP			
	8714149*PED	May 08, 2031				
	8978966	Jan 13, 2032	DP			
	8978966*PED	Jul 13, 2032				
	9216260	Jun 28, 2031	DP			
	9216260*PED	Dec 28, 2031				
	9463288	May 19, 2025	DP			
	9463288*PED	Nov 19, 2025				
	9616024*PED	Mar 01, 2025				
	9731087	May 18, 2031	DP			
	9731087*PED	Nov 18, 2031				
	9782550	Aug 28, 2035	DP			
	9782550*PED	Feb 28, 2036				
	9782551	Aug 28, 2035	DP			
	9782551*PED	Feb 28, 2036				
<u>FLUTICASONE PROPIONATE - XHANCE</u>						
N 209022 001	10076614	Oct 20, 2034	DP		I-940	Mar 15, 2027
	10076615	Jul 30, 2029	U-2133			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>FLUTICASONE PROPIONATE - XHANCE</u>						
N 209022 001	10124132	Mar 06, 2027	DP U-2133			
	10179216	Jul 08, 2033	DP U-2133			
	10252010	Feb 07, 2031	DP			
	10300229	Jul 07, 2035	DP U-2133			
	10478574	Nov 04, 2033	U-2133			
	11033696	May 20, 2033	DP			
	11554229	Feb 23, 2036	U-2133			
	11602603	Oct 27, 2028	DP U-2133			
	12083270	Apr 04, 2031	U-2133			
	7975690	Dec 29, 2025	U-2133			
	8550073	Oct 22, 2029	DP			
	8978647	Dec 06, 2030	DP			
<u>FLUTICASONE PROPIONATE; SALMETEROL XINAFOATE - AIRDUO RESPICLICK</u>						
N 208799 001	10022510	May 18, 2031	DP		M-61	Jul 09, 2024
	10022510*PED	Nov 18, 2031			PED	Jan 09, 2025
	10124131	May 18, 2031	DP			
	10124131*PED	Nov 18, 2031				
	10195375	Feb 14, 2031	DP			
	10195375*PED	Aug 14, 2031				
	10561808	Jan 01, 2032	DP			
	10561808*PED	Jul 01, 2032				
	10765820	May 19, 2025	DP			
	10765820*PED	Nov 19, 2025				
	11969544	Aug 20, 2039	DP			
	11969544*PED	Feb 20, 2040				
	8651103	Mar 26, 2028	DP			
	8651103*PED	Sep 26, 2028				
	8714149	Feb 25, 2032	DP			
	8714149*PED	Aug 25, 2032				
	8978966	Jan 13, 2032	DP			
	8978966*PED	Jul 13, 2032				
	9066957	Oct 06, 2034	DP U-645			
	9066957*PED	Apr 06, 2035				
	9216260	Jun 28, 2031	DP			
	9216260*PED	Dec 28, 2031				
	9415008	Oct 06, 2034	DP U-645			
	9415008*PED	Apr 06, 2035				
	9463288	May 19, 2025	DP			
	9463288*PED	Nov 19, 2025				
	9616024*PED	Mar 01, 2025				
	9731087	May 18, 2031	DP			
	9731087*PED	Nov 18, 2031				
	9987229*PED	Mar 01, 2025				
<u>FLUTICASONE PROPIONATE; SALMETEROL XINAFOATE - AIRDUO RESPICLICK</u>						
N 208799 002	10022510	May 18, 2031	DP			
	10022510*PED	Nov 18, 2031				
	10124131	May 18, 2031	DP			
	10124131*PED	Nov 18, 2031				
	10195375	Feb 14, 2031	DP			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>FLUTICASONE PROPIONATE; SALMETEROL XINAFOATE - AIRDUO RESPICLICK</u>						
N 208799 002	10195375*PED	Aug 14, 2031				
	10561808	Jan 01, 2032	DP			
	10561808*PED	Jul 01, 2032				
	10765820	May 19, 2025	DP			
	10765820*PED	Nov 19, 2025				
	11969544	Aug 20, 2039	DP			
	11969544*PED	Feb 20, 2040				
	8651103	Mar 26, 2028	DP			
	8651103*PED	Sep 26, 2028				
	8714149	Feb 25, 2032	DP			
	8714149*PED	Aug 25, 2032				
	8978966	Jan 13, 2032	DP			
	8978966*PED	Jul 13, 2032				
	9066957	Oct 06, 2034	DP U-645			
	9066957*PED	Apr 06, 2035				
	9216260	Jun 28, 2031	DP			
	9216260*PED	Dec 28, 2031				
	9463288	May 19, 2025	DP			
	9463288*PED	Nov 19, 2025				
	9616024*PED	Mar 01, 2025				
	9731087	May 18, 2031	DP			
	9731087*PED	Nov 18, 2031				
	9987229*PED	Mar 01, 2025				
<u>FLUTICASONE PROPIONATE; SALMETEROL XINAFOATE - AIRDUO RESPICLICK</u>						
N 208799 003	10022510	May 18, 2031	DP			
	10022510*PED	Nov 18, 2031				
	10124131	May 18, 2031	DP			
	10124131*PED	Nov 18, 2031				
	10195375	Feb 14, 2031	DP			
	10195375*PED	Aug 14, 2031				
	10561808	Jan 01, 2032	DP			
	10561808*PED	Jul 01, 2032				
	10765820	May 19, 2025	DP			
	10765820*PED	Nov 19, 2025				
	11969544	Aug 20, 2039	DP			
	11969544*PED	Feb 20, 2040				
	8651103	Mar 26, 2028	DP			
	8651103*PED	Sep 26, 2028				
	8714149	Feb 25, 2032	DP			
	8714149*PED	Aug 25, 2032				
	8978966	Jan 13, 2032	DP			
	8978966*PED	Jul 13, 2032				
	9066957	Oct 06, 2034	DP U-645			
	9066957*PED	Apr 06, 2035				
	9216260	Jun 28, 2031	DP			
	9216260*PED	Dec 28, 2031				
	9463288	May 19, 2025	DP			
	9463288*PED	Nov 19, 2025				
	9616024*PED	Mar 01, 2025				

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>FLUTICASONE PROPIONATE; SALMETEROL XINAFOATE - AIRDUO RESPICLICK</u>						
N 208799 003	9731087	May 18, 2031	DP			
	9731087*PED	Nov 18, 2031				
	9987229*PED	Mar 01, 2025				
<u>FLUTICASONE PROPIONATE; SALMETEROL XINAFOATE - AIRDUO DIGIHALER</u>						
N 208799 004	10022510	May 18, 2031	DP		M-61	Jul 09, 2024
	10022510*PED	Nov 18, 2031			PED	Jan 09, 2025
	10124131	May 18, 2031	DP			
	10124131*PED	Nov 18, 2031				
	10195375	Feb 14, 2031	DP			
	10195375*PED	Aug 14, 2031				
	10561808	Jan 01, 2032	DP			
	10561808*PED	Jul 01, 2032				
	10569034	Aug 16, 2036	DP			
	10569034*PED	Feb 16, 2037				
	10765820	May 19, 2025	DP			
	10765820*PED	Nov 19, 2025				
	10918816	Dec 14, 2035	DP			
	10918816*PED	Jun 14, 2036				
	11000653	Dec 18, 2038	DP			
	11000653*PED	Jun 18, 2039				
	11173259	Jul 06, 2040	DP			
	11173259*PED	Jan 06, 2041				
	11266796	Feb 22, 2041	DP			
	11266796*PED	Aug 22, 2041				
	11344685	Sep 26, 2039	DP			
	11344685*PED	Mar 26, 2040				
	11351317	Feb 10, 2038	DP			
	11351317*PED	Aug 10, 2038				
	11357935	Sep 24, 2038	DP			
	11439777	May 24, 2040	DP			
	11464923	Jun 19, 2040	DP			
	11969544	Aug 20, 2039	DP			
	11969544*PED	Feb 20, 2040				
	8651103	Mar 26, 2028	DP			
	8651103*PED	Sep 26, 2028				
	8714149	Feb 25, 2032	DP			
	8714149*PED	Aug 25, 2032				
	8978966	Jan 13, 2032	DP			
	8978966*PED	Jul 13, 2032				
	9066957	Oct 06, 2034	DP U-645			
	9066957*PED	Apr 06, 2035				
	9216260	Jun 28, 2031	DP			
	9216260*PED	Dec 28, 2031				
	9415008	Oct 06, 2034	DP U-645			
	9415008*PED	Apr 06, 2035				
	9463288	May 19, 2025	DP			
	9463288*PED	Nov 19, 2025				
	9616024*PED	Mar 01, 2025				
	9731087	May 18, 2031	DP			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>FLUTICASONE PROPIONATE; SALMETEROL XINAFOATE - AIRDUO DIGIHALER</u>						
N 208799 004	9731087*PED	Nov 18, 2031				
	9782550	Aug 28, 2035	DP			
	9782550*PED	Feb 28, 2036				
	9782551	Aug 28, 2035	DP			
	9782551*PED	Feb 28, 2036				
	9987229*PED	Mar 01, 2025				
<u>FLUTICASONE PROPIONATE; SALMETEROL XINAFOATE - AIRDUO DIGIHALER</u>						
N 208799 005	10022510	May 18, 2031	DP			
	10022510*PED	Nov 18, 2031				
	10124131	May 18, 2031	DP			
	10124131*PED	Nov 18, 2031				
	10195375	Feb 14, 2031	DP			
	10195375*PED	Aug 14, 2031				
	10561808	Jan 01, 2032	DP			
	10561808*PED	Jul 01, 2032				
	10569034	Aug 16, 2036	DP			
	10569034*PED	Feb 16, 2037				
	10765820	May 19, 2025	DP			
	10765820*PED	Nov 19, 2025				
	10918816	Dec 14, 2035	DP			
	10918816*PED	Jun 14, 2036				
	11000653	Dec 18, 2038	DP			
	11000653*PED	Jun 18, 2039				
	11173259	Jul 06, 2040	DP			
	11173259*PED	Jan 06, 2041				
	11266796	Feb 22, 2041	DP			
	11266796*PED	Aug 22, 2041				
	11344685	Sep 26, 2039	DP			
	11344685*PED	Mar 26, 2040				
	11351317	Feb 10, 2038	DP			
	11351317*PED	Aug 10, 2038				
	11357935	Sep 24, 2038	DP			
	11439777	May 24, 2040	DP			
	11464923	Jun 19, 2040	DP			
	11969544	Aug 20, 2039	DP			
	11969544*PED	Feb 20, 2040				
	8651103	Mar 26, 2028	DP			
	8651103*PED	Sep 26, 2028				
	8714149	Feb 25, 2032	DP			
	8714149*PED	Aug 25, 2032				
	8978966	Jan 13, 2032	DP			
	8978966*PED	Jul 13, 2032				
	9066957	Oct 06, 2034	DP U-645			
	9066957*PED	Apr 06, 2035				
	9216260	Jun 28, 2031	DP			
	9216260*PED	Dec 28, 2031				
	9463288	May 19, 2025	DP			
	9463288*PED	Nov 19, 2025				
	9616024*PED	Mar 01, 2025				

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>FLUTICASONE PROPIONATE; SALMETEROL XINAFOATE - AIRDUO DIGIHALER</u>						
N 208799 005	9731087	May 18, 2031	DP			
	9731087*PED	Nov 18, 2031				
	9782550	Aug 28, 2035	DP			
	9782550*PED	Feb 28, 2036				
	9782551	Aug 28, 2035	DP			
	9782551*PED	Feb 28, 2036				
	9987229*PED	Mar 01, 2025				
<u>FLUTICASONE PROPIONATE; SALMETEROL XINAFOATE - AIRDUO DIGIHALER</u>						
N 208799 006	10022510	May 18, 2031	DP			
	10022510*PED	Nov 18, 2031				
	10124131	May 18, 2031	DP			
	10124131*PED	Nov 18, 2031				
	10195375	Feb 14, 2031	DP			
	10195375*PED	Aug 14, 2031				
	10561808	Jan 01, 2032	DP			
	10561808*PED	Jul 01, 2032				
	10569034	Aug 16, 2036	DP			
	10569034*PED	Feb 16, 2037				
	10765820	May 19, 2025	DP			
	10765820*PED	Nov 19, 2025				
	10918816	Dec 14, 2035	DP			
	10918816*PED	Jun 14, 2036				
	11000653	Dec 18, 2038	DP			
	11000653*PED	Jun 18, 2039				
	11173259	Jul 06, 2040	DP			
	11173259*PED	Jan 06, 2041				
	11266796	Feb 22, 2041	DP			
	11266796*PED	Aug 22, 2041				
	11344685	Sep 26, 2039	DP			
	11344685*PED	Mar 26, 2040				
	11351317	Feb 10, 2038	DP			
	11351317*PED	Aug 10, 2038				
	11357935	Sep 24, 2038	DP			
	11439777	May 24, 2040	DP			
	11464923	Jun 19, 2040	DP			
	11969544	Aug 20, 2039	DP			
	11969544*PED	Feb 20, 2040				
	8651103	Mar 26, 2028	DP			
	8651103*PED	Sep 26, 2028				
	8714149	Feb 25, 2032	DP			
	8714149*PED	Aug 25, 2032				
	8978966	Jan 13, 2032	DP			
	8978966*PED	Jul 13, 2032				
	9066957	Oct 06, 2034	DP U-645			
	9066957*PED	Apr 06, 2035				
	9216260	Jun 28, 2031	DP			
	9216260*PED	Dec 28, 2031				
	9463288	May 19, 2025	DP			
	9463288*PED	Nov 19, 2025				

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>FLUTICASONE PROPIONATE; SALMETEROL XINAFOATE - AIRDUO DIGIHALER</u>						
N 208799	006	9616024*PED	Mar 01, 2025			
		9731087	May 18, 2031	DP		
		9731087*PED	Nov 18, 2031			
		9782550	Aug 28, 2035	DP		
		9782550*PED	Feb 28, 2036			
		9782551	Aug 28, 2035	DP		
		9782551*PED	Feb 28, 2036			
		9987229*PED	Mar 01, 2025			
<u>FOMEPIZOLE - ANTIZOL</u>						
N 020696	001	7553863	Jun 30, 2027	DS DP		
<u>FORMOTEROL FUMARATE; GLYCOPYRROLATE - BEVESPI AEROSPHERE</u>						
N 208294	001	10716753	May 28, 2030	DP U-2889		
		8324266	May 28, 2030		U-2889	
		8703806	May 28, 2030		U-2889	
		8808713	May 28, 2030	DP U-2889		
		8815258	Mar 17, 2031		U-2889	
		9415009	May 28, 2030		U-2889	
		9463161	May 28, 2030	DP U-2889		
<u>FOSAPREPITANT DIMEGLUMINE - EMEND</u>						
N 022023	001				D-186	May 02, 2025
<u>FOSAPREPITANT DIMEGLUMINE - EMEND</u>						
N 022023	002				D-186	May 02, 2025
<u>FOSAPREPITANT DIMEGLUMINE - FOCINVEZ</u>						
N 216686	001	11065265	Jan 11, 2039	DP		
		12042504	Jan 11, 2039	DP		
<u>FOSCARBIDOPA; FOSLEVODOPA - VYALEV</u>						
N 216962	001	10174061	Oct 21, 2035	DP U-4030		
		10730895	Oct 21, 2035	DS		
		9446059	Oct 21, 2035	DS DP U-4030		
<u>FOSDENOPTERIN HYDROBROMIDE - NULIBRY</u>						
N 214018	001	7504095	Jan 31, 2025	DP U-3092	M-286	Oct 27, 2025
					NCE	Feb 26, 2026
					ODE-342	Feb 26, 2028
<u>FOSNETUPITANT CHLORIDE HYDROCHLORIDE; PALONOSETRON HYDROCHLORIDE - AKYNZEO</u>						
N 210493	001	10208073	May 23, 2032		U-2301	
		10624911	Jun 02, 2037	DP		
		10717721	May 23, 2032	DS		
		10828297	Dec 17, 2030		U-2301	
		11312698	May 23, 2032	DS DP		
		11529362	Jun 02, 2037	DP		
		8426450	May 23, 2032	DS DP		
		8895586	May 23, 2032		U-2301	
		9186357	Nov 18, 2030		U-2301	
		9403772	May 23, 2032	DS	U-2301	
		9908907	May 23, 2032	DS DP		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>FOSNETUPITANT CHLORIDE HYDROCHLORIDE; PALONOSETRON HYDROCHLORIDE - AKYNZEO</u>						
N 210493	002 10208073	May 23, 2032			U-2301	
	10624911	Jun 02, 2037	DP			
	10717721	May 23, 2032	DS			
	10828297	Dec 17, 2030			U-2301	
	11312698	May 23, 2032	DS DP			
	11529362	Jun 02, 2037	DP			
	8426450	May 23, 2032	DS DP			
	8895586	May 23, 2032			U-2301	
	9186357	Nov 18, 2030			U-2301	
	9403772	May 23, 2032	DS		U-2301	
	9908907	May 23, 2032	DS DP			
<u>FOSPHENYTOIN SODIUM - SESQUIENT</u>						
N 210864	001 7635773	Mar 13, 2029	DP			
	8410077	Mar 13, 2029	DP			
	9200088	Mar 13, 2029	DP			
	9493582	Feb 27, 2033	DP			
	9750822	Mar 13, 2029	DP			
<u>FOSPHENYTOIN SODIUM - SESQUIENT</u>						
N 210864	002 7635773	Mar 13, 2029	DP			
	8410077	Mar 13, 2029	DP			
	9200088	Mar 13, 2029	DP			
	9493582	Feb 27, 2033	DP			
	9750822	Mar 13, 2029	DP			
<u>FOSTAMATINIB DISODIUM - TAVALISSE</u>						
N 209299	001 7449458	Sep 04, 2031	DS		ODE-174	Apr 17, 2025
	7538108	Mar 28, 2026	DS	U-2294		
	7989448	Jun 12, 2026	DS	U-2294		
	8163902	Jun 17, 2026	DS	U-2294		
	8211889	Jan 19, 2026	DS			
	8263122	Nov 24, 2030	DP			
	8445485	Jun 17, 2026	DP			
	8652492	Nov 06, 2028	DP			
	8771648	Jul 27, 2032	DP			
	8912170	Jun 17, 2026		U-2294		
	8951504	Jul 27, 2032		U-2294		
	9266912	Jan 19, 2026		U-2294		
	9283238	Jun 17, 2026		U-2294		
	RE48898	Jan 19, 2026	DP			
<u>FOSTAMATINIB DISODIUM - TAVALISSE</u>						
N 209299	002 7449458	Sep 04, 2031	DS		ODE-174	Apr 17, 2025
	7538108	Mar 28, 2026	DS	U-2294		
	7989448	Jun 12, 2026	DS	U-2294		
	8163902	Jun 17, 2026	DS	U-2294		
	8211889	Jan 19, 2026	DS			
	8263122	Nov 24, 2030	DP			
	8445485	Jun 17, 2026	DP			
	8652492	Nov 06, 2028	DP			
	8771648	Jul 27, 2032	DP			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>FOSTAMATINIB DISODIUM - TAVALISSE</u>						
N 209299 002	8912170	Jun 17, 2026	U-2294			
	8951504	Jul 27, 2032	U-2294			
	9266912	Jan 19, 2026	U-2294			
	9283238	Jun 17, 2026	U-2294			
	RE48898	Jan 19, 2026	DP			
<u>FOSTEMSAVIR TROMETHAMINE - RUKOBIA</u>						
N 212950 001	7745625	Nov 19, 2027	DS		NCE	Jul 02, 2025
	8168615	Jul 13, 2029	DP			
	8461333	Feb 25, 2025	DS			
<u>FRUQUINTINIB - FRUZAOLA</u>						
N 217564 001	10519142	Sep 07, 2035	DS DP U-3753		NCE	Nov 08, 2028
	11046674	Sep 07, 2035	U-3753			
	7829574	May 09, 2028	DS DP			
	8212033	May 09, 2028	U-3753			
<u>FRUQUINTINIB - FRUZAOLA</u>						
N 217564 002	10519142	Sep 07, 2035	DS DP U-3753		NCE	Nov 08, 2028
	11046674	Sep 07, 2035	U-3753			
	7829574	May 09, 2028	DS DP			
	8212033	May 09, 2028	U-3753			
<u>FULVESTRANT - FULVESTRANT</u>						
N 210326 001	10188663	Feb 14, 2034	DP U-2540			
	9271990	May 17, 2034	DP U-2540			
	9833459	Feb 14, 2034	DP U-2540			
<u>FUROSEMIDE - FUROSCIX</u>						
N 209988 001	10272064	Apr 03, 2034	DP		NP	Oct 07, 2025
	11433044	Apr 03, 2034	U-3462			
	9884039	Apr 03, 2034	U-3462			
<u>FUTIBATINIB - LYTGOBI</u>						
N 214801 001	10434103	Mar 31, 2036	DS DP		NCE	Sep 30, 2027
	11833151	Nov 05, 2039	DP		ODE-410	Sep 30, 2029
	9108973	Feb 23, 2033	DS DP U-3456			
<u>GABAPENTIN ENACARBIL - HORIZANT</u>						
N 022399 001	6818787	Apr 06, 2025	DS DP			
	8026279	Nov 10, 2026	DS DP			
	8114909	Apr 11, 2026	U-1231			
	8686034	Jan 24, 2025	U-1231			
	8686034	Jan 24, 2025	U-1247			
	8795725	Jun 10, 2029	DP U-1231			
	8795725	Jun 10, 2029	DP U-1247			
<u>GABAPENTIN ENACARBIL - HORIZANT</u>						
N 022399 002	6818787	Apr 06, 2025	DS DP			
	8026279	Nov 10, 2026	DS DP			
	8114909	Apr 11, 2026	U-1231			
	8686034	Jan 24, 2025	U-1231			
	8686034	Jan 24, 2025	U-1247			
	8795725	Jun 10, 2029	DP U-1231			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>GABAPENTIN ENACARBIL - HORIZANT</u>						
N 022399 002	8795725	Jun 10, 2029	DP U-1247			
<u>GADOPICLENOL - ELUCIREM</u>						
N 216986 001	10973934	Aug 06, 2039	DS		NCE	Sep 21, 2027
	11590246	Jan 17, 2040	DP			
	8114863	Sep 19, 2028	DS			
<u>GADOPICLENOL - ELUCIREM</u>						
N 216986 002	10973934	Aug 06, 2039	DS		NCE	Sep 21, 2027
	11590246	Jan 17, 2040	DP			
	8114863	Sep 19, 2028	DS			
<u>GADOPICLENOL - ELUCIREM</u>						
N 216986 003	10973934	Aug 06, 2039	DS		NCE	Sep 21, 2027
	11590246	Jan 17, 2040	DP			
	8114863	Sep 19, 2028	DS			
<u>GADOPICLENOL - ELUCIREM</u>						
N 216986 004	10973934	Aug 06, 2039	DS		NCE	Sep 21, 2027
	11590246	Jan 17, 2040	DP			
	8114863	Sep 19, 2028	DS			
<u>GADOPICLENOL - ELUCIREM</u>						
N 216986 005	10973934	Aug 06, 2039	DS		NCE	Sep 21, 2027
	11590246	Jan 17, 2040	DP			
	8114863	Sep 19, 2028	DS			
<u>GADOPICLENOL - ELUCIREM</u>						
N 216986 006	10973934	Aug 06, 2039	DS		NCE	Sep 21, 2027
	11590246	Jan 17, 2040	DP			
	8114863	Sep 19, 2028	DS			
<u>GADOPICLENOL - ELUCIREM</u>						
N 216986 007	10973934	Aug 06, 2039	DS		NCE	Sep 21, 2027
	11590246	Jan 17, 2040	DP			
	8114863	Sep 19, 2028	DS			
<u>GALLIUM DOTATATE GA-68 - NETSPOT</u>						
N 208547 001	9375498	Aug 10, 2032	DP			
<u>GALLIUM GA-68 EDOTREOTIDE - GALLIUM GA 68 EDOTREOTIDE</u>						
N 210828 001					ODE-383	Aug 21, 2026
					W	Aug 21, 2026
<u>GALLIUM GA-68 GOZETOTIDE - GALLIUM GA 68 GOZETOTIDE</u>						
N 212642 001					NCE	Dec 01, 2025
					W	Dec 01, 2025
<u>GALLIUM GA-68 GOZETOTIDE - GALLIUM GA 68 GOZETOTIDE</u>						
N 212643 001					NCE	Dec 01, 2025
					W	Dec 01, 2025
<u>GALLIUM GA-68 GOZETOTIDE - ILLUCCIX</u>						
N 214032 001	11027031	Jul 28, 2035	U-3317			
<u>GALLIUM GA-68 GOZETOTIDE - LOCAMETZ</u>						
N 215841 001	11369590	Aug 15, 2028	DS DP U-3400		NP	Mar 23, 2025
	12109277	Mar 09, 2036	U-4002			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>GALLIUM GA-68 GOZETOTIDE - LOCAMETZ</u>						
N 215841	001 11369590	Aug 15, 2028	DS DP U-3400		NP	Mar 23, 2025
	12109277	Mar 09, 2036	U-4002			
<u>GANAXOLONE - ZTALMY</u>						
N 215904	001 10603308	Aug 10, 2037	U-3374		NCE	Jun 01, 2027
	12144801	Aug 10, 2037	U-4034		ODE-395	Jun 01, 2029
	7858609	Nov 28, 2026	DP			
	8022054	Nov 28, 2026	DP			
	8318714	Nov 28, 2026	DP			
	8367651	Nov 28, 2026	DP			
	8618087	Nov 28, 2026	U-3374			
	9029355	Nov 28, 2026	DP			
	9056116	Nov 28, 2026	U-3374			
<u>GANCICLOVIR - GANZYK-RTU</u>						
N 209347	001 9486530	Sep 02, 2034	DP			
<u>GEMCITABINE HYDROCHLORIDE - INFUGEM</u>						
N 208313	001 9241948	Jul 01, 2033	DP			
<u>GEMCITABINE HYDROCHLORIDE - INFUGEM</u>						
N 208313	002 9241948	Jul 01, 2033	DP			
<u>GEMCITABINE HYDROCHLORIDE - INFUGEM</u>						
N 208313	003 9241948	Jul 01, 2033	DP			
<u>GEMCITABINE HYDROCHLORIDE - INFUGEM</u>						
N 208313	004 9241948	Jul 01, 2033	DP			
<u>GEMCITABINE HYDROCHLORIDE - INFUGEM</u>						
N 208313	005 9241948	Jul 01, 2033	DP			
<u>GEMCITABINE HYDROCHLORIDE - INFUGEM</u>						
N 208313	006 9241948	Jul 01, 2033	DP			
<u>GEMCITABINE HYDROCHLORIDE - INFUGEM</u>						
N 208313	007 9241948	Jul 01, 2033	DP			
<u>GEMCITABINE HYDROCHLORIDE - INFUGEM</u>						
N 208313	008 9241948	Jul 01, 2033	DP			
<u>GEMCITABINE HYDROCHLORIDE - INFUGEM</u>						
N 208313	009 9241948	Jul 01, 2033	DP			
<u>GEMCITABINE HYDROCHLORIDE - INFUGEM</u>						
N 208313	010 9241948	Jul 01, 2033	DP			
<u>GEPIRONE HYDROCHLORIDE - EXXUA</u>						
N 021164	001 7538116	Sep 02, 2025	U-3699		NCE	Sep 22, 2028
<u>GEPIRONE HYDROCHLORIDE - EXXUA</u>						
N 021164	002 7538116	Sep 02, 2025	U-3699		NCE	Sep 22, 2028
<u>GEPIRONE HYDROCHLORIDE - EXXUA</u>						
N 021164	003 7538116	Sep 02, 2025	U-3699		NCE	Sep 22, 2028
<u>GEPIRONE HYDROCHLORIDE - EXXUA</u>						
N 021164	004 7538116	Sep 02, 2025	U-3699		NCE	Sep 22, 2028

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>GILTERITINIB FUMARATE - XOSPATA</u>						
N 211349	001	10786500	Jul 01, 2036	DP	ODE-222	Nov 28, 2025
		11938130	Jul 01, 2036	DP		
		11938131	Jul 01, 2036	DP		
		11938132	Jul 01, 2036	DP		
		11938133	Jul 01, 2036	DP		
		11944620	Jul 01, 2036	DP		
		8969336	Nov 28, 2032	DS DP		
		9487491	Jul 28, 2030	U-2456		
<u>GIVINOSTAT HYDROCHLORIDE - DUVYZAT</u>						
N 217865	001	10688047	Oct 28, 2036	DP U-3885	NCE	Mar 21, 2029
		7329689	Jan 15, 2026	DS DP	ODE-473	Mar 21, 2031
		9421184	Feb 03, 2032	U-3885		
		9867799	Feb 03, 2032	U-3885		
<u>GIVOSIRAN SODIUM - GIVLAARI</u>						
N 212194	001	10119143	Oct 03, 2034	DS DP U-2672	ODE-273	Nov 20, 2026
		10125364	Mar 15, 2033	DS DP U-2672		
		10131907	Aug 24, 2028	DS DP U-2672		
		11028392	Oct 03, 2034	DS DP U-2672		
		8106022	Dec 12, 2029	DS DP U-2672		
		8828956	Dec 04, 2028	DS DP U-2672		
		9133461	Nov 30, 2033	DS DP U-2672		
		9150605	Aug 28, 2025	DS DP		
		9631193	Mar 15, 2033	U-2672		
<u>GLASDEGIB MALEATE - DAURISMO</u>						
N 210656	001	10414748	Apr 13, 2036	DS DP	ODE-224	Nov 21, 2025
		11168066	Apr 13, 2036	U-3254		
		11891372	Apr 13, 2036	U-3254		
		8148401	Jan 30, 2031	DS DP		
		8431597	Jun 29, 2028	DP		
<u>GLASDEGIB MALEATE - DAURISMO</u>						
N 210656	002	10414748	Apr 13, 2036	DS DP	ODE-224	Nov 21, 2025
		11168066	Apr 13, 2036	U-3254		
		11891372	Apr 13, 2036	U-3254		
		8148401	Jan 30, 2031	DS DP		
		8431597	Jun 29, 2028	DP		
<u>GLECAPREVIR; PIBRENTASVIR - MAVYRET</u>						
N 209394	001	10028937	Jun 10, 2030	U-2141	ODE-232	Apr 30, 2026
		10028937	Jun 10, 2030	U-3237	ODE-233	Apr 30, 2026
		10028937*PED	Dec 10, 2030		ODE-372	Jun 10, 2028
		10039754	Jun 10, 2030	U-2141	PED	Oct 30, 2026
		10039754	Jun 10, 2030	U-3237	PED	Oct 30, 2026
		10039754*PED	Dec 10, 2030		PED	Dec 10, 2028
		10286029	Mar 14, 2034	U-3237		
		10286029*PED	Sep 14, 2034			
		11246866	Jun 24, 2036	DP		
		11246866*PED	Dec 24, 2036			
		11484534	Mar 14, 2034	U-3237		
		11484534*PED	Sep 14, 2034			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>GLECAPREVIR; PIBRENTASVIR - MAVYRET</u>						
N 209394	001 8648037	Jan 19, 2032	DS DP U-2141			
	8648037	Jan 19, 2032	DS DP U-3237			
	8648037*PED	Jul 19, 2032				
	8937150	May 18, 2032	DS DP			
	8937150*PED	Nov 18, 2032				
	9321807	Jun 05, 2035	DS			
	9321807*PED	Dec 05, 2035				
	9586978	Nov 06, 2030		U-2141		
	9586978	Nov 06, 2030		U-3237		
	9586978*PED	May 06, 2031				
	RE48923	May 08, 2035	DS			
	RE48923*PED	Nov 08, 2035				
<u>GLECAPREVIR; PIBRENTASVIR - MAVYRET</u>						
N 215110	001 10028937	Jun 10, 2030		U-3238	ODE-372	Jun 10, 2028
	10028937*PED	Dec 10, 2030			PED	Dec 10, 2028
	10039754	Jun 10, 2030		U-3238		
	10039754*PED	Dec 10, 2030				
	10286029	Mar 14, 2034		U-3238		
	10286029*PED	Sep 14, 2034				
	8648037	Jan 19, 2032	DS DP U-3238			
	8648037*PED	Jul 19, 2032				
	8937150	May 18, 2032	DS DP			
	8937150*PED	Nov 18, 2032				
	9321807	Jun 05, 2035	DS DP			
	9321807*PED	Dec 05, 2035				
	9586978	Nov 06, 2030		U-3238		
	9586978*PED	May 06, 2031				
	RE48923	May 08, 2035	DS			
	RE48923*PED	Nov 08, 2035				
<u>GLIMEPIRIDE; PIOGLITAZONE HYDROCHLORIDE - DUETACT</u>						
N 021925	001 7700128	Jan 30, 2027	DP			
	8071130	Jun 08, 2028	DP			
<u>GLIMEPIRIDE; PIOGLITAZONE HYDROCHLORIDE - DUETACT</u>						
N 021925	002 7700128	Jan 30, 2027	DP			
	8071130	Jun 08, 2028	DP			
<u>GLUCAGON - BAOSIMI</u>						
N 210134	001 10213487	Feb 16, 2036	DP U-2604			
	10765602	Sep 23, 2039	DP			
	10894133	Jan 03, 2038	DP			
<u>GLUCAGON - GVOKE PFS</u>						
N 212097	001 11590205	Apr 22, 2036	DP U-2742			
	9649364	Apr 22, 2036	DP U-2742			
<u>GLUCAGON - GVOKE PFS</u>						
N 212097	002 11590205	Apr 22, 2036	DP U-2742			
	9649364	Apr 22, 2036	DP U-2742			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>GLUCAGON - GVOKE HYPOPEN</u>						
N 212097	003	11590205	Apr 22, 2036	DP	U-2742	
	9649364	Apr 22, 2036	DP	U-2742		
<u>GLUCAGON - GVOKE HYPOPEN</u>						
N 212097	004	11590205	Apr 22, 2036	DP	U-2742	
	9649364	Apr 22, 2036	DP	U-2742		
<u>GLUCAGON - GVOKE KIT</u>						
N 212097	005	11590205	Apr 22, 2036	DP	U-2742	
<u>GLYCEROL PHENYL BUTYRATE - RAVICTI</u>						
N 203284	001	10045958	Sep 22, 2030	U-1816		
	10045959	Sep 22, 2030	U-1816			
	10183002	Sep 22, 2030	U-1816			
	10183003	Sep 22, 2030	U-1816			
	10183004	Sep 22, 2030	U-1816			
	10183005	Sep 22, 2030	U-1816			
	10183006	Sep 22, 2030	U-1816			
	10668040	Sep 22, 2030	U-1816			
	8404215	Mar 09, 2032	U-1383	Y		
	8642012	Sep 22, 2030	U-1383			
	9254278	Sep 22, 2030	U-1816			
	9326966	Sep 22, 2030	U-1816			
	9561197	Sep 22, 2030	U-1383			
	9962359	Sep 22, 2030	U-1816			
	9999608	Sep 22, 2030	U-1816			
<u>GLYCOPYRROLATE - SEEBRI</u>						
N 207923	001	8182838	Oct 20, 2028	DP		
	8479730	Oct 11, 2028	DP			
<u>GLYCOPYRROLATE - LONHALA MAGNAIR KIT</u>						
N 208437	001	10376661	Sep 14, 2035	DP		
	10688518	Nov 12, 2036	DP			
	10744277	Dec 07, 2036	DP	U-2941		
	10940110	Feb 26, 2029	DP	U-1773		
	11278683	Aug 16, 2026	DP			
	7931212	Nov 25, 2025	DP			
	9168556	Sep 01, 2032	DP			
	9265900	Dec 07, 2028	DP			
	9604018	May 16, 2033	DP			
	9789270	Oct 30, 2030	DP			
<u>GLYCOPYRROLATE; INDACATEROL MALEATE - UTIBRON</u>						
N 207930	001	6878721	Feb 25, 2025	DS DP	U-1773	
	8182838	Oct 20, 2028	DP			
	8479730	Oct 11, 2028	DP			
<u>GLYCOPYRROLATE; NEOSTIGMINE METHYLSULFATE - PREVDUO</u>						
N 216903	001	10456354	Oct 25, 2038	DP		
	11110054	Oct 25, 2038	DP			
	11938217	Oct 25, 2038	DP			
	12151020	Oct 25, 2038	DP			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>GLYCOPYRRONIUM TOSYLATE - QBREXZA</u>						
N 210361	001 10004717	Feb 28, 2033	DP U-2398			
	10052267	Oct 17, 2028	DP U-2398			
	10543192	Feb 28, 2033	DP			
	10548875	Feb 28, 2033	DS DP U-2398			
	8618160	Dec 10, 2029	DP U-2398			
	8859610	Feb 28, 2033	DP U-2398			
	9259414	Feb 28, 2033	U-2398			
	9744105	Jul 18, 2030	DP U-2398			
<u>GOLODIRSEN - VYONDYS 53</u>						
N 211970	001 10227590	Jun 28, 2025	DS DP		ODE-280	Dec 12, 2026
	10266827	Jun 28, 2025	U-2675			
	10421966	Jun 28, 2025	DS DP			
	10968450	Jun 28, 2025	DS DP			
	10995337	Jun 28, 2025	DP U-2675			
	9024007	Jun 28, 2025	DS DP			
	9994851	Jun 28, 2025	DS DP			
	RE47691	Jun 28, 2028	DP			
<u>GRANISETRON - SANCUSO</u>						
N 022198	001 7608282	Jan 22, 2025	DP U-1011			
<u>GUAIFENESIN; HYDROCODONE BITARTRATE - OBREDON</u>						
N 205474	001 10105324	Nov 13, 2035	DS DP U-2023			
	9549907	Nov 13, 2035	DS DP U-2023			
	9808431	Nov 13, 2035	DS DP U-2023			
<u>HALOBETASOL PROPIONATE - ULTRAVATE</u>						
N 208183	001 8962028	Jun 19, 2033	DP U-1775			
<u>HALOBETASOL PROPIONATE - BRYHALI</u>						
N 209355	001 10478502	Nov 02, 2031	DP U-2625			
	11839656	Nov 02, 2031	U-2625			
	11957753	Nov 02, 2031	DP			
	11986527	Nov 02, 2031	U-2625			
	12076403	Nov 02, 2031	DP			
	8809307	Nov 02, 2031	DP			
<u>HALOBETASOL PROPIONATE - LEXETTE</u>						
N 210566	001 10857159	Nov 30, 2036	DP			
	10857159*PED	May 30, 2037				
	11020407	Nov 30, 2036	DP U-3143			
<u>HALOBETASOL PROPIONATE; TAZAROTENE - DUOBRII</u>						
N 209354	001 10251895	Jun 06, 2036	DP			
	10426787	Jun 06, 2036	U-2625			
	10478502	Nov 02, 2031	DP U-2625			
	11648256	Jun 06, 2036	DP U-2625			
	11679115	Jun 06, 2036	DP U-2625			
	11839656	Nov 02, 2031	U-2625			
	11957753	Nov 02, 2031	DP			
	11986527	Nov 02, 2031	U-2625			
	12076403	Nov 02, 2031	DP			
	8809307	Nov 02, 2031	DP			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>HALOBETASOL PROPIONATE; TAZAROTENE - DUOBRII</u>						
N 209354 001	10251895	Jun 06, 2036	DP			
	10426787	Jun 06, 2036	U-2625			
	10478502	Nov 02, 2031	DP U-2625			
	11648256	Jun 06, 2036	DP U-2625			
	11679115	Jun 06, 2036	DP U-2625			
	11839656	Nov 02, 2031	U-2625			
	11957753	Nov 02, 2031	DP			
	11986527	Nov 02, 2031	U-2625			
	12076403	Nov 02, 2031	DP			
	8809307	Nov 02, 2031	DP			
<u>HEPARIN SODIUM; TAUROLIDINE - DEFENCATH</u>						
N 214520 001	11738120	Apr 15, 2042	DS DP		NCE	Nov 15, 2028
	7696182	May 16, 2025	DS DP U-3774		GAIN	Nov 15, 2033
<u>HEPARIN SODIUM; TAUROLIDINE - DEFENCATH</u>						
N 214520 002	11738120	Apr 15, 2042	DS DP		NCE	Nov 15, 2028
	7696182	May 16, 2025	DS DP U-3774		GAIN	Nov 15, 2033
<u>HEXAMINOLEVULINATE HYDROCHLORIDE - CYSVIEW KIT</u>						
N 022555 001	10556010	Dec 19, 2036	U-2250			
	11235168	Jan 04, 2038	U-2250			
	11311620	Dec 19, 2036	U-2250			
<u>HISTRELIN ACETATE - SUPPRELIN LA</u>						
N 022058 001	8062652	Jun 16, 2026	U-1197			
<u>HYDROCODONE BITARTRATE - ZOHYDRO ER</u>						
N 202880 001	10028946	Jul 25, 2033	U-1810			
	10092559	Sep 12, 2034	U-55			
	10322120	Jul 25, 2033	DP			
	10456393	Jul 25, 2033	U-1810			
	10722511	Jul 25, 2033	U-1810			
	9132096	Sep 12, 2034	DP			
	9265760	Jul 25, 2033	U-1810			
	9326982	Jul 25, 2033	U-1810			
	9333201	Jul 25, 2033	U-1810			
	9339499	Jul 25, 2033	U-1810			
	9421200	Jul 25, 2033	U-1810			
	9433619	Jul 25, 2033	U-1810			
	9452163	Sep 12, 2034	U-55			
	9486451	Sep 12, 2034	U-55			
	9610286	Jul 25, 2033	U-1810			
	9713611	Sep 12, 2034	DP U-55			
<u>HYDROCODONE BITARTRATE - ZOHYDRO ER</u>						
N 202880 002	10028946	Jul 25, 2033	U-1810			
	10092559	Sep 12, 2034	U-55			
	10322120	Jul 25, 2033	DP			
	10456393	Jul 25, 2033	U-1810			
	10722511	Jul 25, 2033	U-1810			
	9132096	Sep 12, 2034	DP			
	9265760	Jul 25, 2033	U-1810			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>HYDROCODONE BITARTRATE - ZOHYDRO ER</u>						
N 202880	002	9326982	Jul 25, 2033	U-1810		
		9333201	Jul 25, 2033	U-1810		
		9339499	Jul 25, 2033	U-1810		
		9421200	Jul 25, 2033	U-1810		
		9433619	Jul 25, 2033	U-1810		
		9452163	Sep 12, 2034	U-55		
		9486451	Sep 12, 2034	U-55		
		9610286	Jul 25, 2033	U-1810		
		9713611	Sep 12, 2034	DP U-55		
<u>HYDROCODONE BITARTRATE - ZOHYDRO ER</u>						
N 202880	003	10028946	Jul 25, 2033	U-1810		
		10092559	Sep 12, 2034	U-55		
		10322120	Jul 25, 2033	DP		
		10456393	Jul 25, 2033	U-1810		
		10722511	Jul 25, 2033	U-1810		
		9132096	Sep 12, 2034	DP		
		9265760	Jul 25, 2033	U-1810		
		9326982	Jul 25, 2033	U-1810		
		9333201	Jul 25, 2033	U-1810		
		9339499	Jul 25, 2033	U-1810		
		9421200	Jul 25, 2033	U-1810		
		9433619	Jul 25, 2033	U-1810		
		9452163	Sep 12, 2034	U-55		
		9486451	Sep 12, 2034	U-55		
		9610286	Jul 25, 2033	U-1810		
		9713611	Sep 12, 2034	DP U-55		
<u>HYDROCODONE BITARTRATE - ZOHYDRO ER</u>						
N 202880	004	10028946	Jul 25, 2033	U-1810		
		10092559	Sep 12, 2034	U-55		
		10322120	Jul 25, 2033	DP		
		10456393	Jul 25, 2033	U-1810		
		10722511	Jul 25, 2033	U-1810		
		9132096	Sep 12, 2034	DP		
		9265760	Jul 25, 2033	U-1810		
		9326982	Jul 25, 2033	U-1810		
		9333201	Jul 25, 2033	U-1810		
		9339499	Jul 25, 2033	U-1810		
		9421200	Jul 25, 2033	U-1810		
		9433619	Jul 25, 2033	U-1810		
		9452163	Sep 12, 2034	U-55		
		9486451	Sep 12, 2034	U-55		
		9610286	Jul 25, 2033	U-1810		
		9713611	Sep 12, 2034	DP U-55		
<u>HYDROCODONE BITARTRATE - ZOHYDRO ER</u>						
N 202880	005	10028946	Jul 25, 2033	U-1810		
		10092559	Sep 12, 2034	U-55		
		10322120	Jul 25, 2033	DP		
		10456393	Jul 25, 2033	U-1810		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>HYDROCODONE BITARTRATE - ZOHYDRO ER</u>						
N 202880	005	10722511	Jul 25, 2033		U-1810	
		9132096	Sep 12, 2034		DP	
		9265760	Jul 25, 2033		U-1810	
		9326982	Jul 25, 2033		U-1810	
		9333201	Jul 25, 2033		U-1810	
		9339499	Jul 25, 2033		U-1810	
		9421200	Jul 25, 2033		U-1810	
		9433619	Jul 25, 2033		U-1810	
		9452163	Sep 12, 2034		U-55	
		9486451	Sep 12, 2034		U-55	
		9610286	Jul 25, 2033		U-1810	
		9713611	Sep 12, 2034		DP U-55	
<u>HYDROCODONE BITARTRATE - ZOHYDRO ER</u>						
N 202880	006	10028946	Jul 25, 2033		U-1810	
		10092559	Sep 12, 2034		U-55	
		10322120	Jul 25, 2033		DP	
		10456393	Jul 25, 2033		U-1810	
		10722511	Jul 25, 2033		U-1810	
		9132096	Sep 12, 2034		DP	
		9265760	Jul 25, 2033		U-1810	
		9326982	Jul 25, 2033		U-1810	
		9333201	Jul 25, 2033		U-1810	
		9339499	Jul 25, 2033		U-1810	
		9421200	Jul 25, 2033		U-1810	
		9433619	Jul 25, 2033		U-1810	
		9452163	Sep 12, 2034		U-55	
		9486451	Sep 12, 2034		U-55	
		9610286	Jul 25, 2033		U-1810	
		9713611	Sep 12, 2034		DP U-55	
<u>HYDROCODONE BITARTRATE - HYSINGLA ER</u>						
N 206627	001	11304908	Aug 24, 2027		DP	
		11304909	Aug 24, 2027		U-1556	
		11844865	Feb 13, 2025		DP U-1556	
		8808740	Dec 21, 2031		DP U-1556	
		9084816	Aug 24, 2027		DP	
		9095614	Aug 24, 2027		U-1556	
		9095615	Aug 24, 2027		DP	
		9486412	Aug 24, 2027		DP	
		9486413	Aug 24, 2027		DP	
		9492389	Aug 24, 2027		DP	
		9492390	Aug 24, 2027		U-1556	
		9492391	Aug 24, 2027		U-1556	
		9545380	Aug 24, 2027		U-1556	
		9572779	Dec 21, 2031		DP	
		9750703	Dec 21, 2031		DP	
		9763933	Aug 24, 2027		DP	
		9770416	Aug 24, 2027		DP	
		9775809	Aug 24, 2027		DP	
		9861584	Dec 21, 2031		DP	

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>HYDROCODONE BITARTRATE - HYSINGLA ER</u>						
N 206627	001 9872837	Dec 21, 2031	DP			
<u>HYDROCODONE BITARTRATE - HYSINGLA ER</u>						
N 206627	002 11304908	Aug 24, 2027	DP			
	11304909	Aug 24, 2027		U-1556		
	11844865	Feb 13, 2025	DP	U-1556		
	8808740	Dec 21, 2031	DP	U-1556		
	9084816	Aug 24, 2027	DP			
	9095614	Aug 24, 2027		U-1556		
	9095615	Aug 24, 2027	DP			
	9486412	Aug 24, 2027	DP			
	9486413	Aug 24, 2027	DP			
	9492389	Aug 24, 2027	DP			
	9492390	Aug 24, 2027		U-1556		
	9492391	Aug 24, 2027		U-1556		
	9545380	Aug 24, 2027		U-1556		
	9572779	Dec 21, 2031	DP			
	9750703	Dec 21, 2031	DP			
	9763933	Aug 24, 2027	DP			
	9770416	Aug 24, 2027	DP			
	9775809	Aug 24, 2027	DP			
	9861584	Dec 21, 2031	DP			
	9872837	Dec 21, 2031	DP			
<u>HYDROCODONE BITARTRATE - HYSINGLA ER</u>						
N 206627	003 11304908	Aug 24, 2027	DP			
	11304909	Aug 24, 2027		U-1556		
	11844865	Feb 13, 2025	DP	U-1556		
	8808740	Dec 21, 2031	DP	U-1556		
	9084816	Aug 24, 2027	DP			
	9095614	Aug 24, 2027		U-1556		
	9095615	Aug 24, 2027	DP			
	9486412	Aug 24, 2027	DP			
	9486413	Aug 24, 2027	DP			
	9492389	Aug 24, 2027	DP			
	9492390	Aug 24, 2027		U-1556		
	9492391	Aug 24, 2027		U-1556		
	9545380	Aug 24, 2027		U-1556		
	9572779	Dec 21, 2031	DP			
	9750703	Dec 21, 2031	DP			
	9763933	Aug 24, 2027	DP			
	9770416	Aug 24, 2027	DP			
	9775809	Aug 24, 2027	DP			
	9861584	Dec 21, 2031	DP			
	9872837	Dec 21, 2031	DP			
<u>HYDROCODONE BITARTRATE - HYSINGLA ER</u>						
N 206627	004 11304908	Aug 24, 2027	DP			
	11304909	Aug 24, 2027		U-1556		
	11844865	Feb 13, 2025	DP	U-1556		
	8808740	Dec 21, 2031	DP	U-1556		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>HYDROCODONE BITARTRATE - HYSINGLA ER</u>						
N 206627	004	9084816	Aug 24, 2027	DP		
		9095614	Aug 24, 2027		U-1556	
		9095615	Aug 24, 2027	DP		
		9486412	Aug 24, 2027	DP		
		9486413	Aug 24, 2027	DP		
		9492389	Aug 24, 2027	DP		
		9492390	Aug 24, 2027		U-1556	
		9492391	Aug 24, 2027		U-1556	
		9545380	Aug 24, 2027		U-1556	
		9572779	Dec 21, 2031	DP		
		9750703	Dec 21, 2031	DP		
		9763933	Aug 24, 2027	DP		
		9770416	Aug 24, 2027	DP		
		9775809	Aug 24, 2027	DP		
		9861584	Dec 21, 2031	DP		
		9872837	Dec 21, 2031	DP		
<u>HYDROCODONE BITARTRATE - HYSINGLA ER</u>						
N 206627	005	11304908	Aug 24, 2027	DP		
		11304909	Aug 24, 2027		U-1556	
		11844865	Feb 13, 2025	DP	U-1556	
		8808740	Dec 21, 2031	DP	U-1556	
		9084816	Aug 24, 2027	DP		
		9095614	Aug 24, 2027		U-1556	
		9095615	Aug 24, 2027	DP		
		9486412	Aug 24, 2027	DP		
		9486413	Aug 24, 2027	DP		
		9492389	Aug 24, 2027	DP		
		9492390	Aug 24, 2027		U-1556	
		9492391	Aug 24, 2027		U-1556	
		9545380	Aug 24, 2027		U-1556	
		9572779	Dec 21, 2031	DP		
		9750703	Dec 21, 2031	DP		
		9763933	Aug 24, 2027	DP		
		9770416	Aug 24, 2027	DP		
		9775809	Aug 24, 2027	DP		
		9861584	Dec 21, 2031	DP		
		9872837	Dec 21, 2031	DP		
<u>HYDROCODONE BITARTRATE - HYSINGLA ER</u>						
N 206627	006	11304908	Aug 24, 2027	DP		
		11304909	Aug 24, 2027		U-1556	
		11844865	Feb 13, 2025	DP	U-1556	
		8808740	Dec 21, 2031	DP	U-1556	
		9084816	Aug 24, 2027	DP		
		9095614	Aug 24, 2027		U-1556	
		9095615	Aug 24, 2027	DP		
		9486412	Aug 24, 2027	DP		
		9486413	Aug 24, 2027	DP		
		9492389	Aug 24, 2027	DP		
		9492390	Aug 24, 2027		U-1556	

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>HYDROCODONE BITARTRATE - HYSINGLA ER</u>						
N 206627	006	9492391	Aug 24, 2027	U-1556		
		9545380	Aug 24, 2027	U-1556		
		9572779	Dec 21, 2031	DP		
		9750703	Dec 21, 2031	DP		
		9763933	Aug 24, 2027	DP		
		9770416	Aug 24, 2027	DP		
		9775809	Aug 24, 2027	DP		
		9861584	Dec 21, 2031	DP		
		9872837	Dec 21, 2031	DP		
<u>HYDROCODONE BITARTRATE - HYSINGLA ER</u>						
N 206627	007	11304908	Aug 24, 2027	DP		
		11304909	Aug 24, 2027	U-1556		
		11844865	Feb 13, 2025	DP U-1556		
		8808740	Dec 21, 2031	DP U-1556		
		9084816	Aug 24, 2027	DP		
		9095614	Aug 24, 2027	U-1556		
		9095615	Aug 24, 2027	DP		
		9486412	Aug 24, 2027	DP		
		9486413	Aug 24, 2027	DP		
		9492389	Aug 24, 2027	DP		
		9492390	Aug 24, 2027	U-1556		
		9492391	Aug 24, 2027	U-1556		
		9545380	Aug 24, 2027	U-1556		
		9572779	Dec 21, 2031	DP		
		9750703	Dec 21, 2031	DP		
		9763933	Aug 24, 2027	DP		
		9770416	Aug 24, 2027	DP		
		9775809	Aug 24, 2027	DP		
		9861584	Dec 21, 2031	DP		
		9872837	Dec 21, 2031	DP		
<u>HYDROCODONE BITARTRATE - VANTRELA ER</u>						
N 207975	001	8445018	Jul 31, 2029	DP		
		9216176	Sep 13, 2027	DP		
		9572803	Sep 13, 2027	DP		
<u>HYDROCODONE BITARTRATE - VANTRELA ER</u>						
N 207975	002	8445018	Jul 31, 2029	DP		
		9216176	Sep 13, 2027	DP		
		9572803	Sep 13, 2027	DP		
<u>HYDROCODONE BITARTRATE - VANTRELA ER</u>						
N 207975	003	8445018	Jul 31, 2029	DP		
		9216176	Sep 13, 2027	DP		
		9572803	Sep 13, 2027	DP		
<u>HYDROCODONE BITARTRATE - VANTRELA ER</u>						
N 207975	004	8445018	Jul 31, 2029	DP		
		9216176	Sep 13, 2027	DP		
		9572803	Sep 13, 2027	DP		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>HYDROCODONE BITARTRATE - VANTRELA ER</u>						
N 207975	005 8445018	Jul 31, 2029	DP			
	9216176	Sep 13, 2027	DP			
	9572803	Sep 13, 2027	DP			
<u>HYDROCORTISONE - ALKINDI SPRINKLE</u>						
N 213876	001 9649280	May 12, 2034	DP U-3075			
	9675559	Jan 10, 2033	U-3075			
	9717740	Nov 19, 2032	U-3075			
<u>HYDROCORTISONE - ALKINDI SPRINKLE</u>						
N 213876	002 9649280	May 12, 2034	DP U-3075			
	9675559	Jan 10, 2033	U-3075			
	9717740	Nov 19, 2032	U-3075			
<u>HYDROCORTISONE - ALKINDI SPRINKLE</u>						
N 213876	003 9649280	May 12, 2034	DP U-3075			
	9675559	Jan 10, 2033	U-3075			
	9717740	Nov 19, 2032	U-3075			
<u>HYDROCORTISONE - ALKINDI SPRINKLE</u>						
N 213876	004 9649280	May 12, 2034	DP U-3075			
	9675559	Jan 10, 2033	U-3075			
	9717740	Nov 19, 2032	U-3075			
<u>HYDROCORTISONE BUTYRATE - LOCOID</u>						
N 022076	001 7981877	Jan 23, 2025	DP			
<u>HYDROCORTISONE SODIUM SUCCINATE - HYDROCORTISONE SODIUM SUCCINATE</u>						
A 214050	001				CGT	Mar 25, 2025
<u>HYDROGEN PEROXIDE - ESKATA</u>						
N 209305	001 10098910	Apr 21, 2035	DP U-2205			
	10493103	Apr 21, 2035	DP			
	10729720	Apr 21, 2035	DP			
	9675639	Jul 04, 2035	DP U-2205			
	9980983	Apr 21, 2035	U-2205			
<u>HYDROMORPHONE HYDROCHLORIDE - HYDROMORPHONE HYDROCHLORIDE</u>						
A 216899	001				PC	Apr 12, 2025
<u>HYDROMORPHONE HYDROCHLORIDE - DILAUDID-HP</u>						
N 019034	001 9248229	Mar 12, 2034	DP			
	9731082	Apr 23, 2032	DP			
<u>HYDROMORPHONE HYDROCHLORIDE - DILAUDID-HP</u>						
N 019034	002 9248229	Mar 12, 2034	DP			
	9731082	Apr 23, 2032	DP			
<u>HYDROMORPHONE HYDROCHLORIDE - DILAUDID</u>						
N 019034	003 9248229	Mar 12, 2034	DP			
	9731082	Apr 23, 2032	DP			
<u>HYDROMORPHONE HYDROCHLORIDE - DILAUDID</u>						
N 019034	004 9248229	Mar 12, 2034	DP			
	9731082	Apr 23, 2032	DP			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>HYDROMORPHONE HYDROCHLORIDE - DILAUDID</u>						
N 019034 005	9248229	Mar 12, 2034	DP			
	9731082	Apr 23, 2032	DP			
<u>HYDROMORPHONE HYDROCHLORIDE - DILAUDID</u>						
N 019034 006	9248229	Mar 12, 2034	DP			
	9731082	Apr 23, 2032	DP			
<u>HYDROMORPHONE HYDROCHLORIDE - DILAUDID</u>						
N 019034 007	9248229	Mar 12, 2034	DP			
	9731082	Apr 23, 2032	DP			
<u>HYDROXYUREA - XROMI</u>						
N 216593 001					NP	Apr 04, 2027
					ODE-470	Apr 04, 2031
<u>IBREXAFUNGERP CITRATE - BREXAFEMME</u>						
N 214900 001	10174074	Jan 19, 2035	DS DP		I-903	Nov 30, 2025
	10370406	Jan 19, 2035		U-3159	NCE	Jun 01, 2026
	10927142	Jan 19, 2035	DS		GAIN	Jun 01, 2031
	11534433	Jun 10, 2039		U-3159		
	11534433	Jun 10, 2039		U-3508		
	8188085	Aug 28, 2030	DS DP	U-3159		
<u>IBRUTINIB - IMBRUVICA</u>						
N 205552 001	10004746	Jun 03, 2031		U-1684	M-14	Aug 24, 2025
	10004746	Jun 03, 2031		U-1946	NPP	Aug 24, 2025
	10004746	Jun 03, 2031		U-2241	ODE-152	Aug 02, 2024
	10004746	Jun 03, 2031		U-2242	ODE-405	Aug 24, 2029
	10004746*PED	Dec 03, 2031			PED	Feb 02, 2025
	10016435	Jun 03, 2031		U-1650	PED	Feb 24, 2026
	10016435*PED	Dec 03, 2031			PED	Feb 24, 2030
	10106548	Jun 03, 2033	DS DP			
	10106548*PED	Dec 03, 2033				
	10125140	Jun 03, 2033	DS DP			
	10125140*PED	Dec 03, 2033				
	10294231	Jun 03, 2033		DP		
	10294231*PED	Dec 03, 2033				
	10294232	Jun 03, 2033		DP		
	10294232*PED	Dec 03, 2033				
	10463668	Oct 24, 2034		U-2654		
	10463668*PED	Apr 24, 2035				
	10478439	Jun 03, 2031		U-1650		
	10478439	Jun 03, 2031		U-1684		
	10478439	Jun 03, 2031		U-1946		
	10478439	Jun 03, 2031		U-2241		
	10478439	Jun 03, 2031		U-2242		
	10478439	Jun 03, 2031		U-2665		
	10478439	Jun 03, 2031		U-3422		
	10478439*PED	Dec 03, 2031				
	10695350	Oct 24, 2034		U-2846		
	10695350*PED	Apr 24, 2035				
	10751342	Jun 03, 2031		U-1491		
	10751342	Jun 03, 2031		U-1946		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>IBRUTINIB - IMBRUVICA</u>						
N 205552 001	10751342	Jun 03, 2031	U-2943			
	10751342	Jun 03, 2031	U-2944			
	10751342*PED	Dec 03, 2031				
	10752634	Jun 03, 2033	DP			
	10752634*PED	Dec 03, 2033				
	10961251	Jun 03, 2033	DP			
	10961251*PED	Dec 03, 2033				
	11672803	Jun 03, 2031	U-1684			
	11672803	Jun 03, 2031	U-1946			
	11672803	Jun 03, 2031	U-2241			
	11672803	Jun 03, 2031	U-2242			
	7514444	Dec 28, 2026	DS DP			
	7514444*PED	Jun 28, 2027				
	8008309	Nov 13, 2027	DS DP			
	8008309*PED	May 13, 2028				
	8476284	Dec 28, 2026	U-1456			
	8476284	Dec 28, 2026	U-1650			
	8476284	Dec 28, 2026	U-1946			
	8476284	Dec 28, 2026	U-1947			
	8476284*PED	Jun 28, 2027				
	8497277	Dec 28, 2026	U-1456			
	8497277	Dec 28, 2026	U-1491			
	8497277	Dec 28, 2026	U-1650			
	8497277	Dec 28, 2026	U-1946			
	8497277	Dec 28, 2026	U-1947			
	8497277	Dec 28, 2026	U-2241			
	8497277	Dec 28, 2026	U-2242			
	8497277	Dec 28, 2026	U-3422			
	8497277*PED	Jun 28, 2027				
	8563563	Apr 26, 2027	U-1491			
	8563563	Apr 26, 2027	U-1650			
	8563563	Apr 26, 2027	U-1946			
	8563563	Apr 26, 2027	U-2219			
	8563563*PED	Oct 26, 2027				
	8697711	Dec 28, 2026	DS DP			
	8697711*PED	Jun 28, 2027				
	8703780	Dec 28, 2026	U-1491			
	8703780*PED	Jun 28, 2027				
	8735403	Dec 28, 2026	DS DP			
	8735403*PED	Jun 28, 2027				
	8754090	Jun 03, 2031	U-1456	Y		
	8754090*PED	Dec 03, 2031				
	8754091	Dec 28, 2026	DP			
	8754091*PED	Jun 28, 2027				
	8952015	Dec 28, 2026	U-1456			
	8952015	Dec 28, 2026	U-1491			
	8952015	Dec 28, 2026	U-1650			
	8952015	Dec 28, 2026	U-1946			
	8952015	Dec 28, 2026	U-1947			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>IBRUTINIB - IMBRUVICA</u>						
N 205552 001	8952015*PED	Jun 28, 2027				
	8957079	Dec 28, 2026	DS DP			
	8957079*PED	Jun 28, 2027				
	8999999	Jun 03, 2031		U-1683		
	8999999	Jun 03, 2031		U-1684		
	8999999*PED	Dec 03, 2031				
	9125889	Jun 03, 2031		U-1745		
	9125889*PED	Dec 03, 2031				
	9181257	Dec 28, 2026	DS DP			
	9181257*PED	Jun 28, 2027				
	9296753	Oct 30, 2033	DS DP			
	9296753*PED	Apr 30, 2034				
	9540382	Aug 18, 2033		U-1456		
	9540382	Aug 18, 2033		U-1650		
	9540382	Aug 18, 2033		U-1684		
	9540382	Aug 18, 2033		U-1946		
	9540382	Aug 18, 2033		U-1947		
	9540382*PED	Feb 18, 2034				
	9713617	Jun 03, 2033	DP			
	9713617*PED	Dec 03, 2033				
	9725455	Jun 03, 2033	DS			
	9725455*PED	Dec 03, 2033				
	9795604	Oct 24, 2034		U-2150		
	9795604	Oct 24, 2034		U-2969		
	9795604	Oct 24, 2034		U-2970		
	9795604	Oct 24, 2034		U-3422		
	9795604*PED	Apr 24, 2035				
	9801881	Jun 03, 2031		U-1491		
	9801881*PED	Dec 03, 2031				
	9801883	Jun 03, 2031		U-2159		
	9801883*PED	Dec 03, 2031				
<u>IBRUTINIB - IMBRUVICA</u>						
N 205552 002	10004746	Jun 03, 2031		U-1684	M-14	Aug 24, 2025
	10004746	Jun 03, 2031		U-1946	NPP	Aug 24, 2025
	10004746	Jun 03, 2031		U-2241	ODE-405	Aug 24, 2029
	10004746	Jun 03, 2031		U-2242	PED	Feb 24, 2026
	10004746*PED	Dec 03, 2031			PED	Feb 24, 2030
	10016435	Jun 03, 2031		U-1650		
	10016435*PED	Dec 03, 2031				
	10106548	Jun 03, 2033	DS DP			
	10106548*PED	Dec 03, 2033				
	10125140	Jun 03, 2033	DS DP			
	10125140*PED	Dec 03, 2033				
	10294231	Jun 03, 2033	DP			
	10294231*PED	Dec 03, 2033				
	10294232	Jun 03, 2033	DP			
	10294232*PED	Dec 03, 2033				
	10463668	Oct 24, 2034		U-2654		
	10463668*PED	Apr 24, 2035				

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>IBRUTINIB - IMBRUVICA</u>						
N 205552 002	10478439	Jun 03, 2031	U-1650			
	10478439	Jun 03, 2031	U-1684			
	10478439	Jun 03, 2031	U-1946			
	10478439	Jun 03, 2031	U-2241			
	10478439	Jun 03, 2031	U-2242			
	10478439	Jun 03, 2031	U-2665			
	10478439	Jun 03, 2031	U-3422			
	10478439*PED	Dec 03, 2031				
	10695350	Oct 24, 2034	U-2846			
	10695350*PED	Apr 24, 2035				
	10751342	Jun 03, 2031	U-1491			
	10751342	Jun 03, 2031	U-1946			
	10751342	Jun 03, 2031	U-2943			
	10751342	Jun 03, 2031	U-2944			
	10751342*PED	Dec 03, 2031				
	10961251	Jun 03, 2033	DP			
	10961251*PED	Dec 03, 2033				
	11672803	Jun 03, 2031	U-1684			
	11672803	Jun 03, 2031	U-1946			
	11672803	Jun 03, 2031	U-2241			
	11672803	Jun 03, 2031	U-2242			
	7514444	Dec 28, 2026	DS DP			
	7514444*PED	Jun 28, 2027				
	8008309	Nov 13, 2027	DS DP			
	8008309*PED	May 13, 2028				
	8476284	Dec 28, 2026	U-1456			
	8476284	Dec 28, 2026	U-1650			
	8476284	Dec 28, 2026	U-1946			
	8476284	Dec 28, 2026	U-1947			
	8476284*PED	Jun 28, 2027				
	8497277	Dec 28, 2026	U-1456			
	8497277	Dec 28, 2026	U-1491			
	8497277	Dec 28, 2026	U-1650			
	8497277	Dec 28, 2026	U-1946			
	8497277	Dec 28, 2026	U-1947			
	8497277	Dec 28, 2026	U-2241			
	8497277	Dec 28, 2026	U-2242			
	8497277	Dec 28, 2026	U-3422			
	8497277*PED	Jun 28, 2027				
	8563563	Apr 26, 2027	U-1491			
	8563563	Apr 26, 2027	U-1650			
	8563563	Apr 26, 2027	U-1946			
	8563563	Apr 26, 2027	U-2219			
	8563563*PED	Oct 26, 2027				
	8697711	Dec 28, 2026	DS DP			
	8697711*PED	Jun 28, 2027				
	8703780	Dec 28, 2026	U-1491			
	8703780*PED	Jun 28, 2027				
	8735403	Dec 28, 2026	DS DP			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>IBRUTINIB - IMBRUVICA</u>						
N 205552 002	8735403*PED	Jun 28, 2027				
	8754090	Jun 03, 2031	U-1456	Y		
	8754090*PED	Dec 03, 2031				
	8754091	Dec 28, 2026	DP			
	8754091*PED	Jun 28, 2027				
	8952015	Dec 28, 2026	U-1456			
	8952015	Dec 28, 2026	U-1491			
	8952015	Dec 28, 2026	U-1650			
	8952015	Dec 28, 2026	U-1946			
	8952015	Dec 28, 2026	U-1947			
	8952015*PED	Jun 28, 2027				
	8957079	Dec 28, 2026	DS DP			
	8957079*PED	Jun 28, 2027				
	8999999	Jun 03, 2031	U-1491			
	8999999	Jun 03, 2031	U-1946			
	8999999	Jun 03, 2031	U-2228			
	8999999*PED	Dec 03, 2031				
	9125889	Jun 03, 2031	U-1650			
	9125889*PED	Dec 03, 2031				
	9181257	Dec 28, 2026	DS			
	9181257*PED	Jun 28, 2027				
	9296753	Oct 30, 2033	DS			
	9296753*PED	Apr 30, 2034				
	9540382	Aug 18, 2033	U-1456			
	9540382	Aug 18, 2033	U-1491			
	9540382	Aug 18, 2033	U-1650			
	9540382	Aug 18, 2033	U-1946			
	9540382	Aug 18, 2033	U-1947			
	9540382*PED	Feb 18, 2034				
	9713617	Jun 03, 2033	DP			
	9713617*PED	Dec 03, 2033				
	9725455	Jun 03, 2033	DS			
	9725455*PED	Dec 03, 2033				
	9795604	Oct 24, 2034	U-2969			
	9795604	Oct 24, 2034	U-2970			
	9795604	Oct 24, 2034	U-3422			
	9795604*PED	Apr 24, 2035				
	9801881	Jun 03, 2031	U-1491			
	9801881*PED	Dec 03, 2031				
	9801883	Jun 03, 2031	U-2159			
	9801883*PED	Dec 03, 2031				
<u>IBRUTINIB - IMBRUVICA</u>						
N 210563 001	10004746	Jun 03, 2031	U-1684		M-14	Aug 24, 2025
	10004746	Jun 03, 2031	U-1946		NPP	Aug 24, 2025
	10004746	Jun 03, 2031	U-2241		ODE-405	Aug 24, 2029
	10004746	Jun 03, 2031	U-2242		PED	Feb 24, 2026
	10004746*PED	Dec 03, 2031			PED	Feb 24, 2030
	10010507	Mar 03, 2036	DP			
	10010507*PED	Sep 03, 2036				

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>IBRUTINIB - IMBRUVICA</u>						
N 210563 001	10016435	Jun 03, 2031				U-1650
	10016435*PED	Dec 03, 2031				
	10106548	Jun 03, 2033	DS DP			
	10106548*PED	Dec 03, 2033				
	10125140	Jun 03, 2033	DS DP			
	10125140*PED	Dec 03, 2033				
	10213386	Mar 03, 2036	DP			
	10213386*PED	Sep 03, 2036				
	10463668	Oct 24, 2034				U-2654
	10463668*PED	Apr 24, 2035				
	10478439	Jun 03, 2031				U-1650
	10478439	Jun 03, 2031				U-1684
	10478439	Jun 03, 2031				U-1946
	10478439	Jun 03, 2031				U-2241
	10478439	Jun 03, 2031				U-2242
	10478439	Jun 03, 2031				U-2665
	10478439	Jun 03, 2031				U-3422
	10478439*PED	Dec 03, 2031				
	10695350	Oct 24, 2034				U-2846
	10695350*PED	Apr 24, 2035				
	10751342	Jun 03, 2031				U-1491
	10751342	Jun 03, 2031				U-1946
	10751342	Jun 03, 2031				U-2943
	10751342	Jun 03, 2031				U-2944
	10751342*PED	Dec 03, 2031				
	10752634	Jun 03, 2033	DP			
	10752634*PED	Dec 03, 2033				
	10828259	Mar 03, 2036	DP			
	10828259*PED	Sep 03, 2036				
	10961251	Jun 03, 2033	DP			
	10961251*PED	Dec 03, 2033				
	11672803	Jun 03, 2031				U-1684
	11672803	Jun 03, 2031				U-1946
	11672803	Jun 03, 2031				U-2241
	11672803	Jun 03, 2031				U-2242
	7514444	Dec 28, 2026	DS DP			
	7514444*PED	Jun 28, 2027				
	8008309	Nov 13, 2027	DS DP			
	8008309*PED	May 13, 2028				
	8476284	Dec 28, 2026				U-1650
	8476284	Dec 28, 2026				U-1946
	8476284	Dec 28, 2026				U-2241
	8476284*PED	Jun 28, 2027				
	8497277	Dec 28, 2026				U-1491
	8497277	Dec 28, 2026				U-1650
	8497277	Dec 28, 2026				U-1946
	8497277	Dec 28, 2026				U-2241
	8497277	Dec 28, 2026				U-2242
	8497277	Dec 28, 2026				U-3422

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>IBRUTINIB - IMBRUVICA</u>						
N 210563 001	8497277*PED	Jun 28, 2027				
	8563563	Apr 26, 2027	U-1491			
	8563563	Apr 26, 2027	U-1650			
	8563563	Apr 26, 2027	U-1946			
	8563563	Apr 26, 2027	U-2241			
	8563563	Apr 26, 2027	U-2242			
	8563563*PED	Oct 26, 2027				
	8697711	Dec 28, 2026	DS DP			
	8697711*PED	Jun 28, 2027				
	8703780	Dec 28, 2026	U-1491			
	8703780	Dec 28, 2026	U-2242			
	8703780*PED	Jun 28, 2027				
	8735403	Dec 28, 2026	DS DP			
	8735403*PED	Jun 28, 2027				
	8754091	Dec 28, 2026	DP			
	8754091*PED	Jun 28, 2027				
	8952015	Dec 28, 2026	U-1491			
	8952015	Dec 28, 2026	U-1650			
	8952015	Dec 28, 2026	U-1946			
	8952015	Dec 28, 2026	U-2241			
	8952015	Dec 28, 2026	U-2242			
	8952015*PED	Jun 28, 2027				
	8957079	Dec 28, 2026	DS DP			
	8957079*PED	Jun 28, 2027				
	8999999	Jun 03, 2031	U-1491			
	8999999	Jun 03, 2031	U-1946			
	8999999	Jun 03, 2031	U-2241			
	8999999	Jun 03, 2031	U-2242			
	8999999*PED	Dec 03, 2031				
	9125889	Jun 03, 2031	U-1650			
	9125889*PED	Dec 03, 2031				
	9181257	Dec 28, 2026	DS			
	9181257*PED	Jun 28, 2027				
	9296753	Oct 30, 2033	DS			
	9296753*PED	Apr 30, 2034				
	9655857	Mar 03, 2036	DP			
	9655857*PED	Sep 03, 2036				
	9725455	Jun 03, 2033	DS			
	9725455*PED	Dec 03, 2033				
	9795604	Oct 24, 2034	U-2969			
	9795604	Oct 24, 2034	U-2970			
	9795604	Oct 24, 2034	U-3422			
	9795604*PED	Apr 24, 2035				
	9801881	Jun 03, 2031	U-1491			
	9801881	Jun 03, 2031	U-2242			
	9801881*PED	Dec 03, 2031				
	9801883	Jun 03, 2031	U-2159			
	9801883	Jun 03, 2031	U-2243			
	9801883*PED	Dec 03, 2031				

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>IBRUTINIB - IMBRUVICA</u>						
N 210563 002	10004746	Jun 03, 2031	U-1684		M-14	Aug 24, 2025
	10004746	Jun 03, 2031	U-1946		NPP	Aug 24, 2025
	10004746	Jun 03, 2031	U-2241		ODE-405	Aug 24, 2029
	10004746	Jun 03, 2031	U-2242		PED	Feb 24, 2026
	10004746*PED	Dec 03, 2031			PED	Feb 24, 2030
	10010507	Mar 03, 2036	DP			
	10010507*PED	Sep 03, 2036				
	10016435	Jun 03, 2031	U-1650			
	10016435*PED	Dec 03, 2031				
	10106548	Jun 03, 2033	DS DP			
	10106548*PED	Dec 03, 2033				
	10125140	Jun 03, 2033	DS DP			
	10125140*PED	Dec 03, 2033				
	10213386	Mar 03, 2036	DP			
	10213386*PED	Sep 03, 2036				
	10463668	Oct 24, 2034	U-2654			
	10463668*PED	Apr 24, 2035				
	10478439	Jun 03, 2031	U-1650			
	10478439	Jun 03, 2031	U-1684			
	10478439	Jun 03, 2031	U-1946			
	10478439	Jun 03, 2031	U-2241			
	10478439	Jun 03, 2031	U-2242			
	10478439	Jun 03, 2031	U-2665			
	10478439	Jun 03, 2031	U-3422			
	10478439*PED	Dec 03, 2031				
	10695350	Oct 24, 2034	U-2846			
	10695350*PED	Apr 24, 2035				
	10751342	Jun 03, 2031	U-1491			
	10751342	Jun 03, 2031	U-1946			
	10751342	Jun 03, 2031	U-2943			
	10751342	Jun 03, 2031	U-2944			
	10751342*PED	Dec 03, 2031				
	10828259	Mar 03, 2036	DP			
	10828259*PED	Sep 03, 2036				
	10961251	Jun 03, 2033	DP			
	10961251*PED	Dec 03, 2033				
	11672803	Jun 03, 2031	U-1684			
	11672803	Jun 03, 2031	U-1946			
	11672803	Jun 03, 2031	U-2241			
	11672803	Jun 03, 2031	U-2242			
	7514444	Dec 28, 2026	DS DP			
	7514444*PED	Jun 28, 2027				
	8008309	Nov 13, 2027	DS DP			
	8008309*PED	May 13, 2028				
	8476284	Dec 28, 2026	U-1650			
	8476284	Dec 28, 2026	U-1946			
	8476284	Dec 28, 2026	U-2241			
	8476284*PED	Jun 28, 2027				
	8497277	Dec 28, 2026	U-1491			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>IBRUTINIB - IMBRUVICA</u>						
N 210563 002	8497277	Dec 28, 2026	U-1650			
	8497277	Dec 28, 2026	U-1946			
	8497277	Dec 28, 2026	U-2241			
	8497277	Dec 28, 2026	U-2242			
	8497277	Dec 28, 2026	U-3422			
	8497277*PED	Jun 28, 2027				
	8563563	Apr 26, 2027	U-1491			
	8563563	Apr 26, 2027	U-1650			
	8563563	Apr 26, 2027	U-1946			
	8563563	Apr 26, 2027	U-2241			
	8563563	Apr 26, 2027	U-2242			
	8563563*PED	Oct 26, 2027				
	8697711	Dec 28, 2026	DS DP			
	8697711*PED	Jun 28, 2027				
	8703780	Dec 28, 2026	U-1491			
	8703780	Dec 28, 2026	U-2242			
	8703780*PED	Jun 28, 2027				
	8735403	Dec 28, 2026	DS DP			
	8735403*PED	Jun 28, 2027				
	8754091	Dec 28, 2026	DP			
	8754091*PED	Jun 28, 2027				
	8952015	Dec 28, 2026	U-1491			
	8952015	Dec 28, 2026	U-1650			
	8952015	Dec 28, 2026	U-1946			
	8952015	Dec 28, 2026	U-2241			
	8952015	Dec 28, 2026	U-2242			
	8952015*PED	Jun 28, 2027				
	8957079	Dec 28, 2026	DS DP			
	8957079*PED	Jun 28, 2027				
	8999999	Jun 03, 2031	U-1491			
	8999999	Jun 03, 2031	U-1946			
	8999999	Jun 03, 2031	U-2241			
	8999999	Jun 03, 2031	U-2242			
	8999999*PED	Dec 03, 2031				
	9125889	Jun 03, 2031	U-1650			
	9125889*PED	Dec 03, 2031				
	9181257	Dec 28, 2026	DS			
	9181257*PED	Jun 28, 2027				
	9296753	Oct 30, 2033	DS			
	9296753*PED	Apr 30, 2034				
	9655857	Mar 03, 2036	DP			
	9655857*PED	Sep 03, 2036				
	9725455	Jun 03, 2033	DS			
	9725455*PED	Dec 03, 2033				
	9795604	Oct 24, 2034	U-2969			
	9795604	Oct 24, 2034	U-2970			
	9795604	Oct 24, 2034	U-3422			
	9795604*PED	Apr 24, 2035				
	9801881	Jun 03, 2031	U-1491			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>IBRUTINIB - IMBRUVICA</u>						
N 210563 002	9801881	Jun 03, 2031	U-2242			
	9801881*PED	Dec 03, 2031				
	9801883	Jun 03, 2031	U-2159			
	9801883	Jun 03, 2031	U-2243			
	9801883*PED	Dec 03, 2031				
<u>IBRUTINIB - IMBRUVICA</u>						
N 210563 003	10004746	Jun 03, 2031	U-1684		M-14	Aug 24, 2025
	10004746	Jun 03, 2031	U-1946		NPP	Aug 24, 2025
	10004746	Jun 03, 2031	U-2241		ODE-405	Aug 24, 2029
	10004746	Jun 03, 2031	U-2242		PED	Feb 24, 2026
	10004746*PED	Dec 03, 2031			PED	Feb 24, 2030
	10010507	Mar 03, 2036	DP			
	10010507*PED	Sep 03, 2036				
	10016435	Jun 03, 2031	U-1650			
	10016435*PED	Dec 03, 2031				
	10106548	Jun 03, 2033	DS DP			
	10106548*PED	Dec 03, 2033				
	10125140	Jun 03, 2033	DS DP			
	10125140*PED	Dec 03, 2033				
	10213386	Mar 03, 2036	DP			
	10213386*PED	Sep 03, 2036				
	10463668	Oct 24, 2034	U-2654			
	10463668*PED	Apr 24, 2035				
	10478439	Jun 03, 2031	U-1650			
	10478439	Jun 03, 2031	U-1684			
	10478439	Jun 03, 2031	U-1946			
	10478439	Jun 03, 2031	U-2241			
	10478439	Jun 03, 2031	U-2242			
	10478439	Jun 03, 2031	U-2665			
	10478439	Jun 03, 2031	U-3422			
	10478439*PED	Dec 03, 2031				
	10695350	Oct 24, 2034	U-2846			
	10695350*PED	Apr 24, 2035				
	10751342	Jun 03, 2031	U-1491			
	10751342	Jun 03, 2031	U-1946			
	10751342	Jun 03, 2031	U-2943			
	10751342	Jun 03, 2031	U-2944			
	10751342*PED	Dec 03, 2031				
	10828259	Mar 03, 2036	DP			
	10828259*PED	Sep 03, 2036				
	10961251	Jun 03, 2033	DP			
	10961251*PED	Dec 03, 2033				
	11672803	Jun 03, 2031	U-1684			
	11672803	Jun 03, 2031	U-1946			
	11672803	Jun 03, 2031	U-2241			
	11672803	Jun 03, 2031	U-2242			
	7514444	Dec 28, 2026	DS DP			
	7514444*PED	Jun 28, 2027				
	8008309	Nov 13, 2027	DS DP			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>IBRUTINIB - IMBRUVICA</u>						
N 210563 003	8008309*PED	May 13, 2028				
	8476284	Dec 28, 2026	U-1650			
	8476284	Dec 28, 2026	U-1946			
	8476284	Dec 28, 2026	U-2241			
	8476284*PED	Jun 28, 2027				
	8497277	Dec 28, 2026	U-1491			
	8497277	Dec 28, 2026	U-1650			
	8497277	Dec 28, 2026	U-1946			
	8497277	Dec 28, 2026	U-2241			
	8497277	Dec 28, 2026	U-2242			
	8497277	Dec 28, 2026	U-3422			
	8497277*PED	Jun 28, 2027				
	8563563	Apr 26, 2027	U-1491			
	8563563	Apr 26, 2027	U-1650			
	8563563	Apr 26, 2027	U-1946			
	8563563	Apr 26, 2027	U-2241			
	8563563	Apr 26, 2027	U-2242			
	8563563*PED	Oct 26, 2027				
	8697711	Dec 28, 2026	DS DP			
	8697711*PED	Jun 28, 2027				
	8703780	Dec 28, 2026	U-1491			
	8703780	Dec 28, 2026	U-2242			
	8703780*PED	Jun 28, 2027				
	8735403	Dec 28, 2026	DS DP			
	8735403*PED	Jun 28, 2027				
	8754091	Dec 28, 2026	DP			
	8754091*PED	Jun 28, 2027				
	8952015	Dec 28, 2026	U-1491			
	8952015	Dec 28, 2026	U-1650			
	8952015	Dec 28, 2026	U-1946			
	8952015	Dec 28, 2026	U-2241			
	8952015	Dec 28, 2026	U-2242			
	8952015*PED	Jun 28, 2027				
	8957079	Dec 28, 2026	DS DP			
	8957079*PED	Jun 28, 2027				
	8999999	Jun 03, 2031	U-1491			
	8999999	Jun 03, 2031	U-1946			
	8999999	Jun 03, 2031	U-2241			
	8999999	Jun 03, 2031	U-2242			
	8999999*PED	Dec 03, 2031				
	9125889	Jun 03, 2031	U-1650			
	9125889*PED	Dec 03, 2031				
	9181257	Dec 28, 2026	DS			
	9181257*PED	Jun 28, 2027				
	9296753	Oct 30, 2033	DS			
	9296753*PED	Apr 30, 2034				
	9655857	Mar 03, 2036	DP			
	9655857*PED	Sep 03, 2036				
	9725455	Jun 03, 2033	DS			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>IBRUTINIB - IMBRUVICA</u>						
N 210563 003	9725455*PED	Dec 03, 2033				
	9795604	Oct 24, 2034	U-2969			
	9795604	Oct 24, 2034	U-2970			
	9795604	Oct 24, 2034	U-3422			
	9795604*PED	Apr 24, 2035				
	9801881	Jun 03, 2031	U-1491			
	9801881	Jun 03, 2031	U-2242			
	9801881*PED	Dec 03, 2031				
	9801883	Jun 03, 2031	U-2159			
	9801883	Jun 03, 2031	U-2243			
	9801883*PED	Dec 03, 2031				
<u>IBRUTINIB - IMBRUVICA</u>						
N 210563 004	10004746	Jun 03, 2031	U-1684		M-14	Aug 24, 2025
	10004746	Jun 03, 2031	U-1946		NPP	Aug 24, 2025
	10004746	Jun 03, 2031	U-2241		ODE-405	Aug 24, 2029
	10004746	Jun 03, 2031	U-2242		PED	Feb 24, 2026
	10004746*PED	Dec 03, 2031			PED	Feb 24, 2030
	10010507	Mar 03, 2036	DP			
	10010507*PED	Sep 03, 2036				
	10016435	Jun 03, 2031	U-1650			
	10016435*PED	Dec 03, 2031				
	10106548	Jun 03, 2033	DS DP			
	10106548*PED	Dec 03, 2033				
	10125140	Jun 03, 2033	DS DP			
	10125140*PED	Dec 03, 2033				
	10213386	Mar 03, 2036	DP			
	10213386*PED	Sep 03, 2036				
	10463668	Oct 24, 2034	U-2654			
	10463668*PED	Apr 24, 2035				
	10478439	Jun 03, 2031	U-1456			
	10478439	Jun 03, 2031	U-1650			
	10478439	Jun 03, 2031	U-1684			
	10478439	Jun 03, 2031	U-1946			
	10478439	Jun 03, 2031	U-1947			
	10478439	Jun 03, 2031	U-2241			
	10478439	Jun 03, 2031	U-2242			
	10478439	Jun 03, 2031	U-2665			
	10478439*PED	Dec 03, 2031				
	10653696	Jun 03, 2031	U-1456			
	10653696*PED	Dec 03, 2031				
	10695350	Oct 24, 2034	U-2846			
	10695350*PED	Apr 24, 2035				
	10751342	Jun 03, 2031	U-1491			
	10751342	Jun 03, 2031	U-1946			
	10751342	Jun 03, 2031	U-2943			
	10751342	Jun 03, 2031	U-2944			
	10751342*PED	Dec 03, 2031				
	10828259	Mar 03, 2036	DP			
	10828259*PED	Sep 03, 2036				

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>IBRUTINIB - IMBRUVICA</u>						
N 210563 004	10961251	Jun 03, 2033	DP			
	10961251*PED	Dec 03, 2033				
	7514444	Dec 28, 2026	DS DP			
	7514444*PED	Jun 28, 2027				
	8008309	Nov 13, 2027	DS DP			
	8008309*PED	May 13, 2028				
	8476284	Dec 28, 2026		U-1456		
	8476284	Dec 28, 2026		U-1650		
	8476284	Dec 28, 2026		U-1946		
	8476284	Dec 28, 2026		U-1947		
	8476284	Dec 28, 2026		U-2241		
	8476284*PED	Jun 28, 2027				
	8497277	Dec 28, 2026		U-1456		
	8497277	Dec 28, 2026		U-1491		
	8497277	Dec 28, 2026		U-1650		
	8497277	Dec 28, 2026		U-1946		
	8497277	Dec 28, 2026		U-1947		
	8497277	Dec 28, 2026		U-2241		
	8497277	Dec 28, 2026		U-2242		
	8497277*PED	Jun 28, 2027				
	8563563	Apr 26, 2027		U-1491		
	8563563	Apr 26, 2027		U-1650		
	8563563	Apr 26, 2027		U-1946		
	8563563	Apr 26, 2027		U-2241		
	8563563	Apr 26, 2027		U-2242		
	8563563*PED	Oct 26, 2027				
	8697711	Dec 28, 2026	DS DP			
	8697711*PED	Jun 28, 2027				
	8703780	Dec 28, 2026		U-1491		
	8703780	Dec 28, 2026		U-2242		
	8703780*PED	Jun 28, 2027				
	8735403	Dec 28, 2026	DS DP			
	8735403*PED	Jun 28, 2027				
	8754090	Jun 03, 2031		U-1456		
	8754090*PED	Dec 03, 2031				
	8754091	Dec 28, 2026	DP			
	8754091*PED	Jun 28, 2027				
	8952015	Dec 28, 2026		U-1456		
	8952015	Dec 28, 2026		U-1491		
	8952015	Dec 28, 2026		U-1650		
	8952015	Dec 28, 2026		U-1946		
	8952015	Dec 28, 2026		U-1947		
	8952015	Dec 28, 2026		U-2241		
	8952015	Dec 28, 2026		U-2242		
	8952015*PED	Jun 28, 2027				
	8957079	Dec 28, 2026	DS DP			
	8957079*PED	Jun 28, 2027				
	8999999	Jun 03, 2031		U-1491		
	8999999	Jun 03, 2031		U-1946		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>IBRUTINIB - IMBRUVICA</u>						
N 210563 004	8999999	Jun 03, 2031	U-2241			
	8999999	Jun 03, 2031	U-2242			
	8999999*PED	Dec 03, 2031				
	9125889	Jun 03, 2031	U-1650			
	9125889*PED	Dec 03, 2031				
	9181257	Dec 28, 2026	DS			
	9181257*PED	Jun 28, 2027				
	9296753	Oct 30, 2033	DS			
	9296753*PED	Apr 30, 2034				
	9655857	Mar 03, 2036	DP			
	9655857*PED	Sep 03, 2036				
	9725455	Jun 03, 2033	DS			
	9725455*PED	Dec 03, 2033				
	9795604	Oct 24, 2034	U-2969			
	9795604	Oct 24, 2034	U-2970			
	9795604*PED	Apr 24, 2035				
	9801881	Jun 03, 2031	U-1491			
	9801881	Jun 03, 2031	U-2242			
	9801881*PED	Dec 03, 2031				
	9801883	Jun 03, 2031	U-2159			
	9801883	Jun 03, 2031	U-2243			
	9801883*PED	Dec 03, 2031				
	9814721	Jun 03, 2031	U-1947			
	9814721*PED	Dec 03, 2031				
<u>IBRUTINIB - IMBRUVICA</u>						
N 217003 001	10004746	Jun 03, 2031	U-3846		NP	Aug 24, 2025
	10004746	Jun 03, 2031	U-3847		ODE-405	Aug 24, 2029
	10004746	Jun 03, 2031	U-3848		PED	Feb 24, 2026
	10004746	Jun 03, 2031	U-3849		PED	Feb 24, 2030
	10016435	Jun 03, 2031	U-3845			
	10106548	Jun 03, 2033	DS DP			
	10106548*PED	Dec 03, 2033				
	10125140	Jun 03, 2033	DS DP			
	10125140*PED	Dec 03, 2033				
	10463668	Oct 24, 2034	U-2970			
	10478439	Jun 03, 2031	U-2666			
	10478439	Jun 03, 2031	U-2970			
	10478439	Jun 03, 2031	U-3422			
	10478439	Jun 03, 2031	U-3842			
	10478439	Jun 03, 2031	U-3843			
	10478439	Jun 03, 2031	U-3844			
	10478439	Jun 03, 2031	U-3845			
	10478439*PED	Dec 03, 2031				
	10695350	Oct 24, 2034	U-2970			
	10751342	Jun 03, 2031	U-2666			
	10751342	Jun 03, 2031	U-3842			
	10751342	Jun 03, 2031	U-3843			
	10751342	Jun 03, 2031	U-3844			
	10961251	Jun 03, 2033	DP			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>IBRUTINIB - IMBRUVICA</u>						
N 217003 001	10961251*PED	Dec 03, 2033				
	11672803	Jun 03, 2031	U-2666			
	11672803	Jun 03, 2031	U-3842			
	11672803	Jun 03, 2031	U-3843			
	11672803	Jun 03, 2031	U-3844			
	7514444	Dec 28, 2026	DS DP			
	7514444*PED	Jun 28, 2027				
	8008309	Nov 13, 2027	DS DP			
	8008309*PED	May 13, 2028				
	8476284	Dec 28, 2026	U-3843			
	8476284	Dec 28, 2026	U-3844			
	8476284	Dec 28, 2026	U-3845			
	8497277	Dec 28, 2026	U-2666			
	8497277	Dec 28, 2026	U-2970			
	8497277	Dec 28, 2026	U-3422			
	8497277	Dec 28, 2026	U-3842			
	8497277	Dec 28, 2026	U-3843			
	8497277	Dec 28, 2026	U-3844			
	8497277	Dec 28, 2026	U-3845			
	8497277*PED	Jun 28, 2027				
	8563563	Apr 26, 2027	U-2666			
	8563563	Apr 26, 2027	U-3842			
	8563563	Apr 26, 2027	U-3843			
	8563563	Apr 26, 2027	U-3844			
	8563563	Apr 26, 2027	U-3845			
	8697711	Dec 28, 2026	DS DP			
	8697711*PED	Jun 28, 2027				
	8703780	Dec 28, 2026	U-2666			
	8703780	Dec 28, 2026	U-3842			
	8735403	Dec 28, 2026	DS DP			
	8735403*PED	Jun 28, 2027				
	8754091	Dec 28, 2026	DP			
	8754091*PED	Jun 28, 2027				
	8952015	Dec 28, 2026	U-2666			
	8952015	Dec 28, 2026	U-3842			
	8952015	Dec 28, 2026	U-3843			
	8952015	Dec 28, 2026	U-3844			
	8952015	Dec 28, 2026	U-3845			
	8957079	Dec 28, 2026	DS DP			
	8957079*PED	Jun 28, 2027				
	8999999	Jun 03, 2031	U-2666			
	8999999	Jun 03, 2031	U-3842			
	8999999	Jun 03, 2031	U-3843			
	8999999	Jun 03, 2031	U-3844			
	9125889	Jun 03, 2031	U-3845			
	9181257	Dec 28, 2026	DS			
	9181257*PED	Jun 28, 2027				
	9296753	Oct 30, 2033	DS			
	9296753*PED	Apr 30, 2034				

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>IBRUTINIB - IMBRUVICA</u>						
N 217003	001 9725455	Jun 03, 2033	DS			
	9725455*PED	Dec 03, 2033				
	9795604	Oct 24, 2034	U-2970			
	9795604	Oct 24, 2034	U-3422			
	9795604*PED	Apr 24, 2035				
	9801881	Jun 03, 2031	U-3846			
	9801881	Jun 03, 2031	U-3847			
	9801883	Jun 03, 2031	U-2666			
	9801883	Jun 03, 2031	U-3842			
	9801883	Jun 03, 2031	U-3843			
	9801883	Jun 03, 2031	U-3844			
<u>IBUPROFEN - CALDOLOR</u>						
N 022348	001				NPP	May 11, 2026
<u>IBUPROFEN - CALDOLOR</u>						
N 022348	002 11806400	Mar 16, 2032	U-3746		NPP	May 11, 2026
	8735452	Sep 30, 2029	U-981			
	8871810	Sep 30, 2029	U-981			
	9012508	Sep 14, 2030	U-981			
	9114068	Sep 30, 2029	U-1735			
	9138404	Sep 30, 2029	U-1756			
	9295639	Sep 30, 2029	U-1756			
	9649284	Sep 30, 2029	U-2018			
<u>IBUPROFEN - CALDOLOR</u>						
N 022348	003 11806400	Mar 16, 2032	U-3746		NPP	May 11, 2026
	8735452	Sep 30, 2029	U-981			
	8871810	Sep 30, 2029	U-981			
	9012508	Sep 14, 2030	U-981			
	9072661	Mar 16, 2032	U-2264			
	9072710	Mar 16, 2032	U-2266			
<u>IBUPROFEN LYSINE - NEOPROFEN</u>						
N 021903	001 8415337	Mar 02, 2032	DS DP			
<u>ICOSAPENT ETHYL - VASCEPA</u>						
N 202057	001 10010517	Apr 29, 2030	U-2690			
	10265287	Apr 29, 2030	U-2700			
	10278935	Jun 28, 2033	U-2701			
	10278936	Jun 28, 2033	U-2702			
	10278937	Jun 28, 2033	U-2703			
	10383840	Jun 28, 2033	U-2704			
	10555924	Jun 28, 2033	U-2743			
	10555925	Jun 28, 2033	U-2744			
	10568861	Jun 28, 2033	U-2756			
	10576054	Jun 28, 2033	U-2762			
	10668042	Jun 28, 2033	U-2841			
	10786478	Jun 28, 2033	U-2959			
	10786478	Jun 28, 2033	U-2960			
	10792267	Apr 29, 2030	U-2961			
	10792270	Jun 28, 2033	U-2962			
	10842766	Apr 29, 2030	U-2997			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>ICOSAPENT ETHYL - VASCEPA</u>						
N 202057 001	10842768	Jun 15, 2030	U-2688			
	10881632	Apr 29, 2030	U-3052			
	10894028	Jun 28, 2033	U-3053			
	11000499	Jun 28, 2033	U-3126			
	11103477	Apr 29, 2030	U-3209			
	11116742	Jun 28, 2033	U-3221			
	11154526	Apr 29, 2030	U-3240			
	11213504	Apr 29, 2030	U-3292			
	11298333	Jun 28, 2033	U-3358			
	11369582	Jun 28, 2033	U-2841			
	11717504	Apr 29, 2030	U-3669			
	8298554	Apr 29, 2030	DP			
	8399446	Feb 09, 2030	U-1287			
	8410086	Jun 15, 2030	U-2688			
	8415335	Feb 09, 2030	U-1287			
	8426399	Feb 09, 2030	U-1287			
	8431560	Feb 09, 2030	U-1287			
	8440650	Feb 09, 2030	U-1287			
	8445003	Apr 29, 2030	U-1287			
	8445013	Apr 29, 2030	U-1287			
	8454994	Apr 29, 2030	U-2689			
	8455472	Jun 15, 2030	U-2690			
	8501225	Apr 29, 2030	U-1287			
	8518929	Feb 09, 2030	U-1287			
	8524698	Feb 09, 2030	U-1287			
	8546372	Feb 09, 2030	U-1287			
	8551521	Apr 29, 2030	U-1287			
	8563608	Apr 29, 2030	U-1287			
	8617593	Apr 29, 2030	U-1478			
	8617593	Apr 29, 2030	U-2691			
	8617594	Apr 29, 2030	U-1287			
	8618166	Apr 29, 2030	U-2689			
	8623406	Apr 29, 2030	U-1478			
	8623406	Apr 29, 2030	U-2692			
	8642077	Apr 29, 2030	U-2693			
	8669245	Jun 15, 2030	U-2694			
	8680144	Feb 09, 2030	U-2695			
	8691871	Apr 29, 2030	U-2689			
	8703185	Apr 29, 2030	U-2691			
	8709475	Apr 29, 2030	U-2689			
	8710041	Jun 15, 2030	U-2690			
	9198892	Sep 25, 2027	U-2706			
	9603826	Jun 28, 2033	U-2696			
	9610272	Jun 28, 2033	U-2697			
	9623001	Jun 28, 2033	U-2698			
	9693984	Jun 28, 2033	U-2697			
	9693985	Jun 28, 2033	U-2696			
	9693986	Jun 28, 2033	U-2698			
	9700537	May 31, 2027	U-2707			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>ICOSAPENT ETHYL - VASCEPA</u>						
N 202057	001 9918954	Jun 28, 2033	U-2699			
<u>ICOSAPENT ETHYL - VASCEPA</u>						
N 202057	002 10010517	Apr 29, 2030	U-2690			
	10265287	Apr 29, 2030	U-2700			
	10278935	Jun 28, 2033	U-2701			
	10278936	Jun 28, 2033	U-2702			
	10278937	Jun 28, 2033	U-2703			
	10383840	Jun 28, 2033	U-2704			
	10555924	Jun 28, 2033	U-2743			
	10555925	Jun 28, 2033	U-2744			
	10568861	Jun 28, 2033	U-2756			
	10576054	Jun 28, 2033	U-2762			
	10668042	Jun 28, 2033	U-2841			
	10786478	Jun 28, 2033	U-2959			
	10786478	Jun 28, 2033	U-2960			
	10792267	Apr 29, 2030	U-2961			
	10792270	Jun 28, 2033	U-2962			
	10842766	Apr 29, 2030	U-2997			
	10842768	Jun 15, 2030	U-2688			
	10881632	Apr 29, 2030	U-3052			
	10894028	Jun 28, 2033	U-3053			
	11000499	Jun 28, 2033	U-3126			
	11103477	Apr 29, 2030	U-3209			
	11116742	Jun 28, 2033	U-3221			
	11154526	Apr 29, 2030	U-3240			
	11213504	Apr 29, 2030	U-3292			
	11298333	Jun 28, 2033	U-3358			
	11369582	Jun 28, 2033	U-2841			
	11717504	Apr 29, 2030	U-3669			
	8298554	Apr 29, 2030	DP			
	8399446	Feb 09, 2030	U-1287			
	8410086	Jun 15, 2030	U-2688			
	8415335	Feb 09, 2030	U-1287			
	8426399	Feb 09, 2030	U-1287			
	8440650	Feb 09, 2030	U-1287			
	8445003	Apr 29, 2030	U-1287			
	8445013	Apr 29, 2030	U-1287			
	8454994	Apr 29, 2030	U-2689			
	8501225	Apr 29, 2030	U-1287			
	8518929	Feb 09, 2030	U-1287			
	8524698	Feb 09, 2030	U-1287			
	8546372	Feb 09, 2030	U-1287			
	8551521	Apr 29, 2030	U-1287			
	8563608	Apr 29, 2030	U-1287			
	8617593	Apr 29, 2030	U-1287			
	8617593	Apr 29, 2030	U-2691			
	8617594	Apr 29, 2030	U-1287			
	8623406	Apr 29, 2030	U-1287			
	8623406	Apr 29, 2030	U-2692			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>ICOSAPENT ETHYL - VASCEPA</u>						
N 202057	002	8642077	Apr 29, 2030	U-2693		
		8669245	Jun 15, 2030	U-2694		
		8680144	Feb 09, 2030	U-2695		
		8691871	Apr 29, 2030	U-2689		
		8703185	Apr 29, 2030	U-2691		
		8709475	Apr 29, 2030	U-2689		
		8710041	Jun 15, 2030	U-2690		
		9198892	Sep 25, 2027	U-2706		
		9603826	Jun 28, 2033	U-2696		
		9610272	Jun 28, 2033	U-2697		
		9623001	Jun 28, 2033	U-2698		
		9693984	Jun 28, 2033	U-2697		
		9693985	Jun 28, 2033	U-2696		
		9693986	Jun 28, 2033	U-2698		
		9700537	May 31, 2027	U-2707		
		9918954	Jun 28, 2033	U-2699		
<u>IDELALISIB - ZYDELIG</u>						
N 205858	001	10730879	Mar 05, 2033	DS DP		
		8865730	Mar 05, 2033	DS DP	U-1615	
		8980901	May 12, 2025		U-1678	
		9149477	May 12, 2025		U-1757	
		9469643	Sep 02, 2033	DS		
		9492449	Mar 11, 2030		U-1914	
		RE44599	Jul 21, 2025		U-1558	
		RE44599	Jul 21, 2025		U-1615	
		RE44638	Aug 05, 2025	DS DP		
<u>IDELALISIB - ZYDELIG</u>						
N 205858	002	10730879	Mar 05, 2033	DS DP		
		8865730	Mar 05, 2033	DS DP	U-1615	
		8980901	May 12, 2025		U-1678	
		9149477	May 12, 2025		U-1757	
		9469643	Sep 02, 2033	DS		
		9492449	Mar 11, 2030		U-1914	
		RE44599	Jul 21, 2025		U-1558	
		RE44599	Jul 21, 2025		U-1615	
		RE44638	Aug 05, 2025	DS DP		
<u>ILOPERIDONE - FANAPT</u>						
N 022192	001	8586610	Nov 02, 2027	U-1625	I-939	Apr 02, 2027
		8652776	Aug 31, 2030	U-1685		
		8999638	Oct 28, 2030	U-1674		
		9072742	Jan 16, 2031	U-1674		
		9074254	Dec 28, 2031	U-1674		
		9074255	Dec 17, 2030	U-1674		
		9074256	Feb 10, 2031	U-1674		
		9138432	Sep 30, 2025	U-1737		
		9157121	Apr 05, 2030	U-1674		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>ILOPERIDONE - FANAPT</u>						
N 022192	002	8586610	Nov 02, 2027	U-1625	I-939	Apr 02, 2027
		8652776	Aug 31, 2030	U-1685		
		8999638	Oct 28, 2030	U-1674		
		9072742	Jan 16, 2031	U-1674		
		9074254	Dec 28, 2031	U-1674		
		9074255	Dec 17, 2030	U-1674		
		9074256	Feb 10, 2031	U-1674		
		9138432	Sep 30, 2025	U-1737		
		9157121	Apr 05, 2030	U-1674		
<u>ILOPERIDONE - FANAPT</u>						
N 022192	003	8586610	Nov 02, 2027	U-1625	I-939	Apr 02, 2027
		8652776	Aug 31, 2030	U-1685		
		8999638	Oct 28, 2030	U-1674		
		9072742	Jan 16, 2031	U-1674		
		9074254	Dec 28, 2031	U-1674		
		9074255	Dec 17, 2030	U-1674		
		9074256	Feb 10, 2031	U-1674		
		9138432	Sep 30, 2025	U-1737		
		9157121	Apr 05, 2030	U-1674		
<u>ILOPERIDONE - FANAPT</u>						
N 022192	004	8586610	Nov 02, 2027	U-1625	I-939	Apr 02, 2027
		8652776	Aug 31, 2030	U-1685		
		8999638	Oct 28, 2030	U-1674		
		9072742	Jan 16, 2031	U-1674		
		9074254	Dec 28, 2031	U-1674		
		9074255	Dec 17, 2030	U-1674		
		9074256	Feb 10, 2031	U-1674		
		9138432	Sep 30, 2025	U-1737		
		9157121	Apr 05, 2030	U-1674		
<u>ILOPERIDONE - FANAPT</u>						
N 022192	005	8586610	Nov 02, 2027	U-1625	I-939	Apr 02, 2027
		8652776	Aug 31, 2030	U-1685		
		8999638	Oct 28, 2030	U-1674		
		9072742	Jan 16, 2031	U-1674		
		9074254	Dec 28, 2031	U-1674		
		9074255	Dec 17, 2030	U-1674		
		9074256	Feb 10, 2031	U-1674		
		9138432	Sep 30, 2025	U-1737		
		9157121	Apr 05, 2030	U-1674		
<u>ILOPERIDONE - FANAPT</u>						
N 022192	006	8586610	Nov 02, 2027	U-1625	I-939	Apr 02, 2027
		8652776	Aug 31, 2030	U-1685		
		8999638	Oct 28, 2030	U-1674		
		9072742	Jan 16, 2031	U-1674		
		9074254	Dec 28, 2031	U-1674		
		9074255	Dec 17, 2030	U-1674		
		9074256	Feb 10, 2031	U-1674		
		9138432	Sep 30, 2025	U-1737		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>ILOPERIDONE - FANAPT</u>						
N 022192	006 9157121	Apr 05, 2030	U-1674			
<u>ILOPERIDONE - FANAPT</u>						
N 022192	007 8586610	Nov 02, 2027	U-1625		I-939	Apr 02, 2027
	8652776	Aug 31, 2030	U-1685			
	8999638	Oct 28, 2030	U-1674			
	9072742	Jan 16, 2031	U-1674			
	9074254	Dec 28, 2031	U-1674			
	9074255	Dec 17, 2030	U-1674			
	9074256	Feb 10, 2031	U-1674			
	9138432	Sep 30, 2025	U-1737			
	9157121	Apr 05, 2030	U-1674			
<u>ILOPROST - AURLUMYN</u>						
N 217933	001				NP	Feb 13, 2027
					ODE-465	Feb 13, 2031
<u>IMATINIB MESYLATE - IMKELDI</u>						
N 219097	001 11957681	Apr 27, 2040	DP			
<u>IMETELSTAT SODIUM - RYTELO</u>						
N 217779	001 7494982	Dec 27, 2025	DS DP		ODE-482	Jun 06, 2031
	9375485	Mar 15, 2033	U-3956			
<u>IMETELSTAT SODIUM - RYTELO</u>						
N 217779	002 7494982	Dec 27, 2025	DS DP		ODE-482	Jun 06, 2031
	9375485	Mar 15, 2033	U-3956			
<u>IMIQUIMOD - ZYCLARA</u>						
N 022483	001 10238644	Dec 11, 2029	U-68			
	10238645	Aug 18, 2029	U-1455			
	10238645	Aug 18, 2029	U-172			
	10918635	Apr 30, 2030	U-1455			
	10918635	Apr 30, 2030	U-172			
	11202752	Apr 30, 2030	U-1455			
	11202752	Apr 30, 2030	U-172			
	11850245	Apr 30, 2030	U-1455			
	11850245	Apr 30, 2030	U-172			
	8236816	Dec 11, 2029	U-68			
	8299109	Dec 11, 2029	U-68			
	8598196	Aug 18, 2029	U-1455			
	8598196	Aug 18, 2029	U-172			
<u>IMIQUIMOD - ZYCLARA</u>						
N 022483	002 11318130	Dec 11, 2029	U-68			
	8222270	Dec 11, 2029	U-68			
<u>INAVOLISIB - ITOVEBI</u>						
N 219249	001 10851091	Jul 01, 2036	DS DP		NCE	Oct 10, 2029
	11028100	Apr 26, 2038	DS DP			
	11760753	Jul 01, 2036	U-4024			
	8242104	Sep 27, 2030	DS DP			
	8343955	Sep 27, 2030	U-4024			
	9650393	Jul 01, 2036	DS DP U-4024			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>INAVOLISIB - ITOVEBI</u>						
N 219249 002	10851091	Jul 01, 2036	DS DP		NCE	Oct 10, 2029
	11028100	Apr 26, 2038	DS DP			
	11760753	Jul 01, 2036		U-4024		
	8242104	Sep 27, 2030	DS DP			
	8343955	Sep 27, 2030		U-4024		
	9650393	Jul 01, 2036	DS DP	U-4024		
<u>INCLISIRAN SODIUM - LEOVIO</u>						
N 214012 001	10125369	Aug 18, 2034	DS DP	U-3652	NCE	Dec 22, 2026
	10131907	Aug 24, 2028	DS DP	U-3652		
	10806791	Dec 04, 2028	DS			
	10851377	Aug 25, 2036		U-3652		
	8106022	Dec 12, 2029	DS DP	U-3652		
	8222222	Dec 29, 2027		U-3652		
	8809292	May 10, 2027	DS DP	U-3652		
	8828956	Dec 04, 2028	DS DP	U-3652		
	9370582	Dec 04, 2028	DS DP	U-3652		
<u>INDACATEROL MALEATE - ARCAPTA NEOHALER</u>						
N 022383 001	6878721	Feb 25, 2025	DS DP	U-1168		
	8479730	Oct 11, 2028	DP			
<u>INDIGOTINDISULFONATE SODIUM - BLUDIGO</u>						
N 216264 001	10927258	Dec 23, 2037	DS		NCE	Jul 08, 2027
	11499050	Dec 23, 2037	DS			
	11845867	Nov 25, 2036	DS			
<u>INDOCYANINE GREEN - SPY AGENT GREEN KIT</u>						
N 211580 001	10631746	Aug 04, 2035	DP	U-2815	I-911	Jun 05, 2026
	11712320	Jul 14, 2039	DP			
	8185176	Jun 04, 2028		U-2462		
	8406860	Apr 09, 2029		U-2463		
	8647605	Feb 11, 2029		U-2464		
	8647605	Feb 11, 2029		U-2468		
	9421280	Nov 24, 2025		U-2466		
	9421280	Nov 24, 2025		U-2467		
<u>INDOMETHACIN - TIVORBEX</u>						
N 204768 001	8734847	Apr 23, 2030	DP			
	8992982	Apr 23, 2030	DP			
	9089471	Apr 23, 2030		U-55		
<u>INDOMETHACIN - TIVORBEX</u>						
N 204768 002	8734847	Apr 23, 2030	DP			
	8992982	Apr 23, 2030	DP			
	9089471	Apr 23, 2030		U-55		
<u>INFIGRATINIB PHOSPHATE - TRUSELTIQ</u>						
N 214622 001	10278969	Dec 11, 2034	DP		NCE	May 28, 2026
	11160804	Dec 11, 2034	DP		ODE-353	May 28, 2028
	8552002	Aug 25, 2029	DS DP			
	9067896	Aug 06, 2028	DS			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>INFIGRATINIB PHOSPHATE - TRUSELTIQ</u>						
N 214622 002	10278969	Dec 11, 2034	DP		NCE	May 28, 2026
	11160804	Dec 11, 2034	DP		ODE-353	May 28, 2028
	8552002	Aug 25, 2029	DS DP			
	9067896	Aug 06, 2028	DS			
<u>INGENOL MEBUTATE - PICATO</u>						
N 202833 001	8278292	Jul 06, 2027	DP			
	8372827	Dec 18, 2026	DP			
	8372828	Dec 18, 2026	DP			
	8377919	Dec 18, 2026	DP			
	8536163	Dec 18, 2026	U-1440			
	8716271	Dec 18, 2026	U-1440			
	8735375	Dec 18, 2026	U-1440			
	9789078	May 15, 2033	U-2138			
	9820959	Dec 18, 2026	DP U-1440			
	9833428	Dec 18, 2026	DP			
	9833429	Dec 18, 2026	DP			
	9861603	Dec 18, 2026	U-1440			
<u>INGENOL MEBUTATE - PICATO</u>						
N 202833 002	8278292	Jul 06, 2027	DP			
	8372827	Dec 18, 2026	DP			
	8372828	Dec 18, 2026	DP			
	8377919	Dec 18, 2026	DP			
	8536163	Dec 18, 2026	U-1440			
	8716271	Dec 18, 2026	U-1440			
	8735375	Dec 18, 2026	U-1440			
	9820959	Dec 18, 2026	DP U-1440			
	9833428	Dec 18, 2026	DP			
	9833429	Dec 18, 2026	DP			
	9861603	Dec 18, 2026	U-1440			
<u>INOTERSEN SODIUM - TEGSEDI</u>						
N 211172 001	8101743	Apr 01, 2025	DS DP		ODE-212	Oct 05, 2025
	8697860	Apr 29, 2031	DP			
	9061044	Apr 29, 2031	DS			
	9399774	Apr 29, 2031	U-2430			
<u>IOBENGUANE I-131 - AZEDRA</u>						
N 209607 001					ODE-204	Jul 30, 2025
<u>IPRATROPIUM BROMIDE - ATROVENT HFA</u>						
N 021527 001	8474447	Jan 17, 2030	DP			
<u>IPTACOPAN HYDROCHLORIDE - FABHALTA</u>						
N 218276 001	10093663	Jul 14, 2034	U-3980		I-949	Aug 07, 2027
	11603363	May 25, 2041	DS DP		NCE	Dec 05, 2028
	11723901	Aug 30, 2038	U-3980		ODE-456	Dec 05, 2030
	11951101	Jul 15, 2041	U-3895			
	9682968	Jul 14, 2034	DS DP			
<u>IRINOTECAN HYDROCHLORIDE - ONIVYDE</u>						
N 207793 001	10456360	Oct 15, 2036	DP		I-932	Feb 13, 2027
	10722508	May 02, 2025	DS DP		ODE-463	Feb 13, 2031

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>IRINOTECAN HYDROCHLORIDE - ONIVYDE</u>						
N 207793 001	10980795	Jun 12, 2033	U-1848			
	10993914	Oct 15, 2036	DP			
	11344552	Aug 19, 2036	U-3824			
	11369597	Jun 12, 2033	U-1848			
	12059497	Oct 15, 2036	DP			
	8147867	Aug 29, 2028	DS DP			
	8329213	Jan 06, 2027	DS DP			
	8703181	May 02, 2025	U-1434			
	8992970	May 02, 2025	DS DP			
	9339497	Jun 12, 2033	U-1848			
	9364473	Jun 12, 2033	U-1856			
	9452162	Jun 12, 2033	U-1899			
	9492442	Jun 12, 2033	U-1848			
	9492442	Jun 12, 2033	U-1899			
	9492442	Jun 12, 2033	U-1917			
	9717724	Jun 12, 2033	U-1848			
	9717724	Jun 12, 2033	U-2091			
	9724303	May 02, 2025	DS DP			
	9730891	May 02, 2025	U-1848			
	9782349	May 02, 2025	DS DP			
<u>ISAVUCONAZONIUM SULFATE - CRESEMBA</u>						
N 207500 001	10206879	Sep 14, 2027	DP		NCE	Mar 06, 2020
	10206879*PED	Mar 14, 2028			NPP	Dec 08, 2026
	10603280	Sep 14, 2027	DP		ODE-305	Mar 06, 2022
	10603280*PED	Mar 14, 2028			ODE-454	Dec 08, 2030
	6812238	Oct 31, 2025	DS		ODE-458	Dec 08, 2030
	6812238*PED	Apr 30, 2026			ODE-90	Mar 06, 2022
					PED	Sep 06, 2020
					PED	Sep 06, 2022
					PED	Sep 06, 2022
					PED	Jun 08, 2027
					PED	Jun 08, 2031
					PED	Jun 08, 2031
					GAIN	Sep 06, 2025
					GAIN	Sep 06, 2027
					GAIN	Sep 06, 2027
<u>ISAVUCONAZONIUM SULFATE - CRESEMBA</u>						
N 207500 002	10206879	Sep 14, 2027	DP		NCE	Mar 06, 2020
	10206879*PED	Mar 14, 2028			NPP	Dec 08, 2026
	10603280	Sep 14, 2027	DP		ODE*	Mar 06, 2022
	10603280*PED	Mar 14, 2028			ODE-454	Dec 08, 2030
	6812238	Oct 31, 2025	DS		ODE-458	Dec 08, 2030
	6812238*PED	Apr 30, 2026			PED	Sep 06, 2020
					PED	Sep 06, 2022
					PED	Jun 08, 2027
					PED	Jun 08, 2031
					PED	Jun 08, 2031
					GAIN	Sep 06, 2025
					GAIN	Sep 06, 2027

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>ISAVUCONAZONIUM SULFATE - CRESEMBA</u>						
N 207500 002	10206879	Sep 14, 2027	DP		NCE	Mar 06, 2020
	10206879*PED	Mar 14, 2028			NPP	Dec 08, 2026
	10603280	Sep 14, 2027	DP		ODE*	Mar 06, 2022
	10603280*PED	Mar 14, 2028			ODE-454	Dec 08, 2030
	6812238	Oct 31, 2025	DS		ODE-458	Dec 08, 2030
	6812238*PED	Apr 30, 2026			PED	Sep 06, 2020
					PED	Sep 06, 2022
					PED	Jun 08, 2027
					PED	Jun 08, 2031
					PED	Jun 08, 2031
					GAIN	Sep 06, 2025
					GAIN	Sep 06, 2027
<u>ISAVUCONAZONIUM SULFATE - CRESEMBA</u>						
N 207501 001	6812238	Oct 31, 2025	DS		NCE	Mar 06, 2020
	6812238*PED	Apr 30, 2026			NPP	Dec 08, 2026
					ODE-305	Mar 06, 2022
					ODE-453	Dec 08, 2030
					ODE-459	Dec 08, 2030
					ODE-90	Mar 06, 2022
					PED	Sep 06, 2020
					PED	Sep 06, 2022
					PED	Sep 06, 2022
					PED	Jun 08, 2027
					PED	Jun 08, 2031
					PED	Jun 08, 2031
					GAIN	Sep 06, 2025
					GAIN	Sep 06, 2027
					GAIN	Sep 06, 2027
<u>ISOPROPYL ALCOHOL - ZURAGARD</u>						
N 210872 001	10688291	Dec 20, 2034	DP U-1397			
	8226971	May 06, 2025	DP			
	8389583	Aug 09, 2029		U-1397		
	8703828	May 23, 2028	DP			
	9011897	Feb 08, 2025	DP			
	9629368	May 23, 2028		U-1397		
	9844654	Apr 24, 2036	DP U-1397			
<u>ISOTRETINOIN - ABSORICA LD</u>						
N 211913 001	9700535	Aug 04, 2035	DP			
	9750711	May 29, 2035	DP			
<u>ISOTRETINOIN - ABSORICA LD</u>						
N 211913 002	9700535	Aug 04, 2035	DP			
	9750711	May 29, 2035	DP			
<u>ISOTRETINOIN - ABSORICA LD</u>						
N 211913 003	9700535	Aug 04, 2035	DP			
	9750711	May 29, 2035	DP			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>ISOTRETINOIN - ABSORICA LD</u>						
N 211913	004 9700535	Aug 04, 2035	DP			
	9750711	May 29, 2035	DP			
<u>ISOTRETINOIN - ABSORICA LD</u>						
N 211913	005 9700535	Aug 04, 2035	DP			
	9750711	May 29, 2035	DP			
<u>ISOTRETINOIN - ABSORICA LD</u>						
N 211913	006 9700535	Aug 04, 2035	DP			
	9750711	May 29, 2035	DP			
<u>ISTRADEFYLLINE - NOURIANZ</u>						
N 022075	001 7727993	Jan 28, 2028	DP U-2623			
	8318201	Sep 05, 2027	DP			
<u>ISTRADEFYLLINE - NOURIANZ</u>						
N 022075	002 7727993	Jan 28, 2028	DP U-2623			
	8318201	Sep 05, 2027	DP			
<u>ITRACONAZOLE - ONMEL</u>						
N 022484	001 8486456	Oct 03, 2028	DP U-1054			
<u>ITRACONAZOLE - TOLSURA</u>						
N 208901	001 10463740	Jun 21, 2033	DP U-2453			
	10806792	Jun 21, 2033	DP			
	8921374	Jun 21, 2033	DP			
	9272046	Jun 21, 2033	DP			
	9713642	Jun 21, 2033	U-2453			
<u>IVABRADINE - CORLANOR</u>						
N 209964	001 7361649	Feb 22, 2026	DS DP U-1694		ODE-234	Apr 22, 2026
	7361650	Feb 22, 2026	DS DP U-1694		PED	Oct 22, 2026
	7867996	Dec 12, 2026	DS DP U-1694			
	7879842	Feb 22, 2026	DS DP U-1694			
<u>IVABRADINE HYDROCHLORIDE - CORLANOR</u>						
N 206143	001 7361649	Feb 22, 2026	DS DP U-1694			
	7361649*PED	Aug 22, 2026				
	7361650	Feb 22, 2026	DS DP U-1694			
	7361650*PED	Aug 22, 2026				
	7867996	Dec 12, 2026	DS DP U-1694			
	7867996*PED	Jun 12, 2027				
	7879842	Feb 22, 2026	DS DP U-1694			
	7879842*PED	Aug 22, 2026				
<u>IVABRADINE HYDROCHLORIDE - CORLANOR</u>						
N 206143	002 7361649	Feb 22, 2026	DS DP U-1694			
	7361649*PED	Aug 22, 2026				
	7361650	Feb 22, 2026	DS DP U-1694			
	7361650*PED	Aug 22, 2026				
	7867996	Dec 12, 2026	DS DP U-1694			
	7867996*PED	Jun 12, 2027				
	7879842	Feb 22, 2026	DS DP U-1694			
	7879842*PED	Aug 22, 2026				

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>IVACAFTOR - KALYDECO</u>						
N 203188	001	10646481	Aug 13, 2029	DP	ODE-199	Aug 15, 2025
		11564916	Aug 13, 2029	U-3530	ODE-338	Dec 21, 2027
		7495103	May 20, 2027	DS DP		
		8324242	Aug 05, 2027	U-1311		
		8324242	Aug 05, 2027	U-1906		
		8354427	Jul 06, 2026	U-1311		
		8354427	Jul 06, 2026	U-1905		
		8410274	Dec 28, 2026	DP		
		8629162	Jun 24, 2025	U-2234		
		8754224	Dec 28, 2026	DS DP		
		9670163	Dec 28, 2026	DP U-1311		
<u>IVACAFTOR - KALYDECO</u>						
N 207925	001	10272046	Feb 27, 2033	DP U-2531	ODE-199	Aug 15, 2025
		10646481	Aug 13, 2029	DP	ODE-236	Apr 29, 2026
		11147770	Feb 27, 2033	DP U-3339	ODE-338	Dec 21, 2027
		11564916	Aug 13, 2029	U-3528	ODE-435	May 03, 2030
		11752106	Feb 27, 2033	DP U-3697		
		7495103	May 20, 2027	DS DP		
		8324242	Aug 05, 2027	U-1311		
		8324242	Aug 05, 2027	U-1906		
		8324242	Aug 05, 2027	U-2527		
		8354427	Jul 06, 2026	U-1311		
		8354427	Jul 06, 2026	U-1905		
		8354427	Jul 06, 2026	U-2528		
		8410274	Dec 28, 2026	DP		
		8629162	Jun 24, 2025	U-2234		
		8629162	Jun 24, 2025	U-2529		
		8754224	Dec 28, 2026	DS DP		
		8883206	Feb 27, 2033	DP		
		9670163	Dec 28, 2026	DP U-1311		
		9670163	Dec 28, 2026	DP U-2530		
<u>IVACAFTOR - KALYDECO</u>						
N 207925	002	10272046	Feb 27, 2033	DP U-2531	ODE-199	Aug 15, 2025
		10646481	Aug 13, 2029	DP	ODE-236	Apr 29, 2026
		11147770	Feb 27, 2033	DP U-3339	ODE-338	Dec 21, 2027
		11564916	Aug 13, 2029	U-3528	ODE-435	May 03, 2030
		11752106	Feb 27, 2033	DP U-3697		
		7495103	May 20, 2027	DS DP		
		8324242	Aug 05, 2027	U-1311		
		8324242	Aug 05, 2027	U-1906		
		8324242	Aug 05, 2027	U-2527		
		8354427	Jul 06, 2026	U-1311		
		8354427	Jul 06, 2026	U-1905		
		8354427	Jul 06, 2026	U-2528		
		8410274	Dec 28, 2026	DP		
		8629162	Jun 24, 2025	U-2234		
		8629162	Jun 24, 2025	U-2529		
		8754224	Dec 28, 2026	DS DP		
		8883206	Feb 27, 2033	DP		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>IVACAFTOR - KALYDECO</u>						
N 207925 002	9670163	Dec 28, 2026	DP U-1311			
	9670163	Dec 28, 2026	DP U-2530			
<u>IVACAFTOR - KALYDECO</u>						
N 207925 003	10272046	Feb 27, 2033	DP U-2967		ODE-199	Aug 15, 2025
	10646481	Aug 13, 2029	DP		ODE-236	Apr 29, 2026
	11147770	Feb 27, 2033	DP U-3339		ODE-338	Dec 21, 2027
	11564916	Aug 13, 2029	U-3528		ODE-435	May 03, 2030
	11752106	Feb 27, 2033	DP U-3697			
	7495103	May 20, 2027	DS DP			
	8324242	Aug 05, 2027	U-1311			
	8324242	Aug 05, 2027	U-1906			
	8324242	Aug 05, 2027	U-2963			
	8354427	Jul 06, 2026	U-1311			
	8354427	Jul 06, 2026	U-1905			
	8354427	Jul 06, 2026	U-2964			
	8410274	Dec 28, 2026	DP			
	8629162	Jun 24, 2025	U-2234			
	8629162	Jun 24, 2025	U-2965			
	8754224	Dec 28, 2026	DS DP			
	8883206	Feb 27, 2033	DP			
	9670163	Dec 28, 2026	DP U-1311			
	9670163	Dec 28, 2026	DP U-2966			
<u>IVACAFTOR - KALYDECO</u>						
N 207925 004	10272046	Feb 27, 2033	DP U-3605		NPP	May 03, 2026
	10646481	Aug 13, 2029	DP		ODE-435	May 03, 2030
	11147770	Feb 27, 2033	DP U-3604			
	11564916	Aug 13, 2029	U-3603			
	11752106	Feb 27, 2033	DP U-3697			
	7495103	May 20, 2027	DS DP			
	8324242	Aug 05, 2027	U-3609			
	8354427	Jul 06, 2026	U-3608			
	8410274	Dec 28, 2026	DP			
	8629162	Jun 24, 2025	U-3607			
	8754224	Dec 28, 2026	DS DP			
	8883206	Feb 27, 2033	DP			
	9670163	Dec 28, 2026	DP U-3606			
<u>IVACAFTOR - KALYDECO</u>						
N 207925 005	10272046	Feb 27, 2033	DP U-3605		NPP	May 03, 2026
	10646481	Aug 13, 2029	DP		ODE-435	May 03, 2030
	11147770	Feb 27, 2033	DP U-3604			
	11564916	Aug 13, 2029	U-3603			
	11752106	Feb 27, 2033	DP U-3697			
	7495103	May 20, 2027	DS DP			
	8324242	Aug 05, 2027	U-3609			
	8354427	Jul 06, 2026	U-3608			
	8410274	Dec 28, 2026	DP			
	8629162	Jun 24, 2025	U-3607			
	8754224	Dec 28, 2026	DS DP			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>IVACAFTOR - KALYDECO</u>						
N 207925	005 8883206	Feb 27, 2033	DP			
	9670163	Dec 28, 2026	DP U-3606			
<u>IVACAFTOR; IVACAFTOR, TEZACAFTOR - SYMDEKO (COPACKAGED)</u>						
N 210491	001 10022352	Apr 09, 2027	DP U-2343		ODE-173	Feb 12, 2025
	10022352	Apr 09, 2027	DP U-2573		ODE-247	Jun 21, 2026
	10058546	Jul 15, 2033	U-3022		ODE-335	Dec 21, 2027
	10058546	Jul 15, 2033	U-3023			
	10081621	Mar 25, 2031	DP U-3024			
	10081621	Mar 25, 2031	DP U-3025			
	10206877	Apr 14, 2035	DP U-3026			
	10206877	Apr 14, 2035	DP U-3027			
	10239867	Apr 09, 2027	DS DP U-2512			
	10239867	Apr 09, 2027	DS DP U-2569			
	10646481	Aug 13, 2029	DP			
	11564916	Aug 13, 2029	U-3527			
	11578062	Mar 25, 2031	DP U-3545			
	11639347	Apr 09, 2027	DS DP U-2569			
	11951212	Apr 14, 2035	DP U-3894			
	7495103	May 20, 2027	DS DP			
	7645789	May 01, 2027	DS DP			
	7776905	Jun 03, 2027	DS DP			
	8324242	Aug 05, 2027	U-2246			
	8354427	Jul 06, 2026	U-3021			
	8410274	Dec 28, 2026	DP			
	8415387	Nov 12, 2027	U-2246			
	8598181	May 01, 2027	U-2246			
	8623905	May 01, 2027	DS DP			
	8629162	Jun 24, 2025	U-2247			
	8754224	Dec 28, 2026	DS DP			
	9012496	Jul 15, 2033	U-2248			
	9670163	Dec 28, 2026	DP U-2246			
	9931334	Dec 28, 2026	DP U-2275			
	9931334	Dec 28, 2026	DP U-2575			
	9974781	Apr 09, 2027	DP U-2318			
	9974781	Apr 09, 2027	DP U-2574			
<u>IVACAFTOR; IVACAFTOR, TEZACAFTOR - SYMDEKO (COPACKAGED)</u>						
N 210491	002 10022352	Apr 09, 2027	DP U-2343		ODE-173	Feb 12, 2025
	10022352	Apr 09, 2027	DP U-2573		ODE-247	Jun 21, 2026
	10058546	Jul 15, 2033	U-3022		ODE-335	Dec 21, 2027
	10058546	Jul 15, 2033	U-3023			
	10081621	Mar 25, 2031	DP U-3024			
	10081621	Mar 25, 2031	DP U-3025			
	10206877	Apr 14, 2035	DP U-3026			
	10206877	Apr 14, 2035	DP U-3027			
	10239867	Apr 09, 2027	DS DP U-2512			
	10239867	Apr 09, 2027	DS DP U-2569			
	10646481	Aug 13, 2029	DP			
	11564916	Aug 13, 2029	U-3527			
	11578062	Mar 25, 2031	DP U-3545			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>IVACAFTOR; IVACAFTOR, TEZACAFTOR - SYMDEKO (COPACKAGED)</u>						
N 210491	002	11639347	Apr 09, 2027	DS DP	U-2569	
		7495103	May 20, 2027	DS DP		
		7645789	May 01, 2027	DS DP		
		7776905	Jun 03, 2027	DS DP		
		8324242	Aug 05, 2027		U-2246	
		8354427	Jul 06, 2026		U-3021	
		8410274	Dec 28, 2026	DP		
		8415387	Nov 12, 2027		U-2246	
		8598181	May 01, 2027		U-2246	
		8623905	May 01, 2027	DS DP		
		8629162	Jun 24, 2025		U-2247	
		8754224	Dec 28, 2026	DS DP		
		9012496	Jul 15, 2033		U-2248	
		9670163	Dec 28, 2026	DP	U-2246	
		9931334	Dec 28, 2026	DP	U-2275	
		9931334	Dec 28, 2026	DP	U-2575	
		9974781	Apr 09, 2027	DP	U-2318	
		9974781	Apr 09, 2027	DP	U-2574	
<u>IVACAFTOR; LUMACAFTOR - ORKAMBI</u>						
N 206038	001	10076513	Dec 04, 2028	DP	U-2411	
		10597384	Dec 04, 2028	DS DP	U-2777	
		10646481	Aug 13, 2029	DP		
		11052075	Dec 04, 2028	DP	U-3181	
		11564916	Aug 13, 2029		U-3529	
		12065432	Dec 04, 2028		U-3984	
		7495103	May 20, 2027	DS DP		
		7973038	Nov 08, 2026		U-1973	
		8324242	Aug 05, 2027		U-1311	
		8324242	Aug 05, 2027		U-1911	
		8410274	Dec 28, 2026	DP		
		8507534	Sep 20, 2030	DS DP		
		8653103	Dec 04, 2028	DP		
		8716338	Sep 20, 2030	DP	U-1718	
		8716338	Sep 20, 2030	DP	U-1910	
		8741933	Nov 08, 2026		U-1717	
		8741933	Nov 08, 2026		U-1909	
		8754224	Dec 28, 2026	DS DP		
		8846718	Jul 02, 2029		U-1717	
		8846718	Jul 02, 2029		U-1908	
		8993600	Dec 11, 2030	DP		
		9150552	Dec 04, 2028		U-1908	
		9192606	Sep 29, 2029	DP	U-1912	
		9216969	Nov 08, 2026	DS DP		
		9670163	Dec 28, 2026	DP	U-1911	
		9931334	Dec 28, 2026	DP	U-2276	
<u>IVACAFTOR; LUMACAFTOR - ORKAMBI</u>						
N 206038	002	10597384	Dec 04, 2028	DS DP	U-2777	
		10646481	Aug 13, 2029	DP		
		11052075	Dec 04, 2028	DP	U-3181	

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>IVACAFTOR; LUMACAFTOR - ORKAMBI</u>						
N 206038	002	11564916	Aug 13, 2029	U-3529		
		12065432	Dec 04, 2028	U-3984		
		7495103	May 20, 2027	DS DP		
		7973038	Nov 08, 2026	U-1973		
		8324242	Aug 05, 2027	U-1911		
		8410274	Dec 28, 2026	DP		
		8507534	Sep 20, 2030	DS DP		
		8653103	Dec 04, 2028	DP		
		8716338	Sep 20, 2030	DP U-1910		
		8741933	Nov 08, 2026	U-1909		
		8754224	Dec 28, 2026	DS DP		
		8846718	Jul 02, 2029	U-1908		
		8993600	Dec 11, 2030	DP		
		9150552	Dec 04, 2028	U-1908		
		9192606	Sep 29, 2029	DP U-1912		
		9216969	Nov 08, 2026	DP		
		9670163	Dec 28, 2026	DP U-1911		
		9931334	Dec 28, 2026	DP U-2276		
<u>IVACAFTOR; LUMACAFTOR - ORKAMBI</u>						
N 211358	001	10597384	Dec 04, 2028	DS DP U-2778	NPP	Sep 02, 2025
		10646481	Aug 13, 2029	DP	ODE-195	Aug 07, 2025
		11564916	Aug 13, 2029	U-3526	ODE-408	Sep 02, 2029
		12065432	Dec 04, 2028	U-3983		
		7495103	May 20, 2027	DS DP		
		7973038	Nov 08, 2026	U-2374		
		8324242	Aug 05, 2027	U-2374		
		8410274	Dec 28, 2026	DP		
		8507534	Sep 20, 2030	DS DP		
		8653103	Dec 04, 2028	DP		
		8716338	Sep 20, 2030	DP U-2396		
		8741933	Nov 08, 2026	U-2374		
		8754224	Dec 28, 2026	DS DP		
		8846718	Jul 02, 2029	U-2375		
		8993600	Dec 11, 2030	DP		
		9150552	Dec 04, 2028	U-2375		
		9192606	Sep 29, 2029	DP U-2397		
		9216969	Nov 08, 2026	DP		
		9670163	Dec 28, 2026	DP U-2376		
		9931334	Dec 28, 2026	DP U-2376		
<u>IVACAFTOR; LUMACAFTOR - ORKAMBI</u>						
N 211358	002	10597384	Dec 04, 2028	DS DP U-2778	NPP	Sep 02, 2025
		10646481	Aug 13, 2029	DP	ODE-195	Aug 07, 2025
		11564916	Aug 13, 2029	U-3526	ODE-408	Sep 02, 2029
		12065432	Dec 04, 2028	U-3983		
		7495103	May 20, 2027	DS DP		
		7973038	Nov 08, 2026	U-2374		
		8324242	Aug 05, 2027	U-2374		
		8410274	Dec 28, 2026	DP		
		8507534	Sep 20, 2030	DS DP		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>IVACAFTOR; LUMACAFTOR - ORKAMBI</u>						
N 211358	002	8653103	Dec 04, 2028	DP		
		8716338	Sep 20, 2030	DP U-2396		
		8741933	Nov 08, 2026	U-2374		
		8754224	Dec 28, 2026	DS DP		
		8846718	Jul 02, 2029	U-2375		
		8993600	Dec 11, 2030	DP		
		9150552	Dec 04, 2028	U-2375		
		9192606	Sep 29, 2029	DP U-2397		
		9216969	Nov 08, 2026	DP		
		9670163	Dec 28, 2026	DP U-2376		
		9931334	Dec 28, 2026	DP U-2376		
<u>IVACAFTOR; LUMACAFTOR - ORKAMBI</u>						
N 211358	003	10597384	Dec 04, 2028	DS DP U-3430	NS	Sep 02, 2025
		10646481	Aug 13, 2029	DP	ODE-408	Sep 02, 2029
		11564916	Aug 13, 2029	U-3526		
		12065432	Dec 04, 2028	U-3983		
		7495103	May 20, 2027	DS DP		
		7973038	Nov 08, 2026	U-3424		
		8324242	Aug 05, 2027	U-3424		
		8410274	Dec 28, 2026	DP		
		8507534	Sep 20, 2030	DS DP		
		8653103	Dec 04, 2028	DP		
		8716338	Sep 20, 2030	DP U-3426		
		8741933	Nov 08, 2026	U-3424		
		8754224	Dec 28, 2026	DS DP		
		8846718	Jul 02, 2029	U-3427		
		8993600	Dec 11, 2030	DP		
		9150552	Dec 04, 2028	U-3427		
		9192606	Sep 29, 2029	DP U-3428		
		9216969	Nov 08, 2026	DP		
		9670163	Dec 28, 2026	DP U-3429		
		9931334	Dec 28, 2026	DP U-3429		
<u>IVERMECTIN - SOOLANTRA</u>						
N 206255	001	10206939	Mar 13, 2034	U-1631		
		9089587	Mar 13, 2034	U-1631		
		9233117	Mar 13, 2034	U-1631		
		9233118	Mar 13, 2034	U-1631		
		9782425	Mar 13, 2034	U-1631		
<u>IVOSIDENIB - TIBSOVO</u>						
N 211192	001	10449184	Mar 13, 2035	DP	I-893	May 25, 2025
		10610125	Jun 21, 2030	U-2784	I-924	Oct 24, 2026
		10610125	Jun 21, 2030	U-2785	ODE-203	Jul 20, 2025
		10610125	Jun 21, 2030	U-3385	ODE-242	May 02, 2026
		10653710	Oct 18, 2036	U-3387	ODE-368	Aug 25, 2028
		10717764	Jan 18, 2033	U-3215	ODE-447	Oct 24, 2030
		10799490	Mar 13, 2035	DP U-2981		
		10799490	Mar 13, 2035	DP U-2982		
		10799490	Mar 13, 2035	DP U-3384		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>IVOSIDENIB - TIBSOVO</u>						
N 211192	001	10980788				
		Jun 07, 2039				U-3112
		10980788				U-3113
		10980788				U-3214
		10980788				U-3383
		10980788				U-3743
		11667673				U-3742
		Jan 18, 2033				U-3742
		9474779		DS DP		U-2350
		Aug 19, 2033		DS DP		U-2533
		9474779		DS DP		U-2534
		Aug 19, 2033		DS DP		U-2534
		9474779		DS DP		U-3213
		Aug 19, 2033		DS DP		U-3213
		9474779		DS DP		U-3386
		Aug 19, 2033		DS DP		U-3386
		9474779		DS DP		U-3742
		Aug 19, 2033		DS DP		U-3742
		9850277		DS DP		U-2350
		Jan 18, 2033		DS DP		U-2350
		9850277		DS DP		U-2533
		Jan 18, 2033		DS DP		U-2533
		9850277		DS DP		U-2534
		Jan 18, 2033		DS DP		U-2534
		9850277		DS DP		U-3213
		Jan 18, 2033		DS DP		U-3213
		9850277		DS DP		U-3386
		Jan 18, 2033		DS DP		U-3386
		9850277		DS DP		U-3742
		Jan 18, 2033		DS DP		U-3742
		9968595		DP		U-2351
		Mar 13, 2035		DP		U-2351
		9968595		DP		U-2533
		Mar 13, 2035		DP		U-2533
		9968595		DP		U-2534
		Mar 13, 2035		DP		U-2534
		9968595		DP		U-3384
		Mar 13, 2035		DP		U-3384
<u>IXAZOMIB CITRATE - NINLARO</u>						
N 208462	001	7442830				
		Nov 20, 2029		DS DP		U-2434
		7687662		DS DP		
		Aug 06, 2027		DS DP		
		8003819		DS DP		U-2434
		Aug 06, 2027		DS DP		U-2434
		8530694		DS DP		U-2434
		Aug 06, 2027		DS DP		U-2434
		8859504		DS DP		
		Jun 16, 2029		DS DP		
		8871745				U-2434
		Aug 06, 2027				U-2434
		9175017				U-2434
		Jun 16, 2029				U-2434
<u>IXAZOMIB CITRATE - NINLARO</u>						
N 208462	002	7442830				
		Nov 20, 2029		DS DP		U-2434
		7687662		DS DP		
		Aug 06, 2027		DS DP		
		8003819		DS DP		U-2434
		Aug 06, 2027		DS DP		U-2434
		8530694		DS DP		U-2434
		Aug 06, 2027		DS DP		U-2434
		8859504		DS DP		
		Jun 16, 2029		DS DP		
		8871745				U-2434
		Aug 06, 2027				U-2434
		9175017				U-2434
		Jun 16, 2029				U-2434
<u>IXAZOMIB CITRATE - NINLARO</u>						
N 208462	003	7442830				
		Nov 20, 2029		DS DP		U-2434
		7687662		DS DP		
		Aug 06, 2027		DS DP		
		8003819		DS DP		U-2434
		Aug 06, 2027		DS DP		U-2434
		8530694		DS DP		U-2434
		Aug 06, 2027		DS DP		U-2434
		8859504		DS DP		
		Jun 16, 2029		DS DP		
		8871745				U-2434
		Aug 06, 2027				U-2434
		9175017				U-2434
		Jun 16, 2029				U-2434

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>KETOROLAC TROMETHAMINE - ACULAR LS</u>						
N 021528	001 8008338	May 24, 2027	DS DP U-1181			
<u>KETOROLAC TROMETHAMINE - ACUVAIL</u>						
N 022427	001 7842714	Aug 15, 2029	DS DP			
	8512717	Mar 07, 2028	DP			
	9192571	Mar 07, 2028	DP			
<u>KETOROLAC TROMETHAMINE; PHENYLEPHRINE HYDROCHLORIDE - OMIIDRIA</u>						
N 205388	001 9066856	Oct 23, 2033	DP			
	9066856*PED	Apr 23, 2034				
	9486406	Oct 23, 2033	DP			
	9486406*PED	Apr 23, 2034				
	9855246	Oct 23, 2033	DP			
<u>KETOTIFEN FUMARATE - ACUVUE THERAVISION WITH KETOTIFEN</u>						
N 022388	001 9474746	Mar 27, 2028	DP		NP	Feb 25, 2025
	9962376	Jun 27, 2030	DP			
<u>LACOSAMIDE - VIMPAT</u>						
N 022253	001				D-188	Apr 28, 2026
<u>LACOSAMIDE - VIMPAT</u>						
N 022253	002				D-188	Apr 28, 2026
<u>LACOSAMIDE - VIMPAT</u>						
N 022253	003				D-188	Apr 28, 2026
<u>LACOSAMIDE - VIMPAT</u>						
N 022253	004				D-188	Apr 28, 2026
<u>LACOSAMIDE - VIMPAT</u>						
N 022254	001				D-188	Apr 28, 2026
<u>LACOSAMIDE - VIMPAT</u>						
N 022255	001				D-188	Apr 28, 2026
<u>LACOSAMIDE - MOTPOLY XR</u>						
N 216185	001 11337943	Jun 05, 2040	DP U-3660			
	11337943	Jun 05, 2040	DP U-3954			
	11883374	Jun 05, 2040	DP U-3660			
	11883374	Jun 05, 2040	DP U-3954			
	12042474	Jun 05, 2040	DP U-3660			
	12042474	Jun 05, 2040	DP U-3954			
<u>LACOSAMIDE - MOTPOLY XR</u>						
N 216185	002 11337943	Jun 05, 2040	DP U-3660			
	11337943	Jun 05, 2040	DP U-3954			
	11883374	Jun 05, 2040	DP U-3660			
	11883374	Jun 05, 2040	DP U-3954			
	12042474	Jun 05, 2040	DP U-3660			
	12042474	Jun 05, 2040	DP U-3954			
<u>LACOSAMIDE - MOTPOLY XR</u>						
N 216185	003 11337943	Jun 05, 2040	DP U-3660			
	11337943	Jun 05, 2040	DP U-3954			
	11883374	Jun 05, 2040	DP U-3660			
	11883374	Jun 05, 2040	DP U-3954			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>LACOSAMIDE - MOTPOLY XR</u>						
N 216185	003 12042474	Jun 05, 2040	DP U-3660			
	12042474	Jun 05, 2040	DP U-3954			
<u>LACTITOL - PIZENSY</u>						
N 211281	001 10806743	May 12, 2037	U-1516		NCE	Feb 12, 2025
<u>LAMIVUDINE; RALTEGRAVIR POTASSIUM - DUTREBIS</u>						
N 206510	001 7754731	Mar 11, 2029	DS DP U-1663			
	7754731*PED	Sep 11, 2029				
<u>LAMOTRIGINE - LAMICTAL XR</u>						
N 022115	001 8637512	Jun 14, 2028	DP			
<u>LAMOTRIGINE - LAMICTAL XR</u>						
N 022115	002 8637512	Jun 14, 2028	DP			
<u>LAMOTRIGINE - LAMICTAL XR</u>						
N 022115	003 8637512	Jun 14, 2028	DP			
<u>LAMOTRIGINE - LAMICTAL XR</u>						
N 022115	004 8637512	Jun 14, 2028	DP			
<u>LAMOTRIGINE - LAMICTAL XR</u>						
N 022115	005 8637512	Jun 14, 2028	DP			
<u>LAMOTRIGINE - LAMICTAL XR</u>						
N 022115	006 8637512	Jun 14, 2028	DP			
<u>LANDIOLOL HYDROCHLORIDE - RAPIBLYK</u>						
N 217202	001				NCE	Nov 22, 2029
<u>LANSOPRAZOLE - LANSOPRAZOLE</u>						
N 208025	001 11077055	Apr 21, 2036	DP			
	11986554	Apr 21, 2036	DP			
<u>LANTHANUM CARBONATE - FOSRENOL</u>						
N 204734	001 8980327	Dec 01, 2030	DP			
	9023397	Dec 01, 2030	DP			
<u>LANTHANUM CARBONATE - FOSRENOL</u>						
N 204734	002 8980327	Dec 01, 2030	DP			
	9023397	Dec 01, 2030	DP			
<u>LAPATINIB DITOSYLATE - TYKERB</u>						
N 022059	001 8821927	Sep 18, 2029	DS DP			
<u>LAROTRECTINIB SULFATE - VITRAKVI</u>						
N 210861	001 10005783	Oct 21, 2029	U-2472		ODE-215	Nov 26, 2025
	10047097	Oct 21, 2029	U-2474		ODE-220	Nov 26, 2025
	10172861	Nov 16, 2035	DS DP		ODE-221	Nov 26, 2025
	10285993	Nov 16, 2035	U-2470			
	10774085	Oct 21, 2029	U-2470			
	10799505	Aug 15, 2036	DS DP			
	10813936	Nov 16, 2035	U-2987			
	8513263	Dec 23, 2029	DS DP			
	8865698	Oct 21, 2029	U-2469			
	9127013	Oct 21, 2029	DS DP			
	9447104	Oct 21, 2029	U-2470			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>LAROTRECTINIB SULFATE - VITRAKVI</u>						
N 210861	001	9676783	Oct 21, 2029	U-2469		
		9782414	Nov 16, 2035	U-2475		
<u>LAROTRECTINIB SULFATE - VITRAKVI</u>						
N 210861	002	10005783	Oct 21, 2029	U-2472	ODE-215	Nov 26, 2025
		10047097	Oct 21, 2029	U-2474	ODE-220	Nov 26, 2025
		10172861	Nov 16, 2035	DS DP	ODE-221	Nov 26, 2025
		10285993	Nov 16, 2035	U-2470		
		10774085	Oct 21, 2029	U-2470		
		10799505	Aug 15, 2036	DS DP		
		10813936	Nov 16, 2035	U-2987		
		8513263	Dec 23, 2029	DS DP		
		8865698	Oct 21, 2029	U-2469		
		9127013	Oct 21, 2029	DS DP		
		9447104	Oct 21, 2029	U-2470		
		9676783	Oct 21, 2029	U-2469		
		9782414	Nov 16, 2035	U-2475		
<u>LAROTRECTINIB SULFATE - VITRAKVI</u>						
N 211710	001	10005783	Oct 21, 2029	U-2472	ODE-215	Nov 26, 2025
		10045991	Apr 04, 2037	U-2473	ODE-220	Nov 26, 2025
		10047097	Oct 21, 2029	U-2474	ODE-221	Nov 26, 2025
		10137127	Apr 04, 2037	DP		
		10172861	Nov 16, 2035	DS		
		10668072	Apr 04, 2037	DP		
		10774085	Oct 21, 2029	U-2470		
		10799505	Aug 15, 2036	DS		
		11191766	Apr 04, 2037	U-2471		
		11484535	Apr 04, 2037	U-2470		
		8513263	Dec 23, 2029	DS DP		
		8865698	Oct 21, 2029	U-2469		
		9127013	Oct 21, 2029	DS DP		
		9447104	Oct 21, 2029	U-2470		
		9676783	Oct 21, 2029	U-2469		
		9782414	Nov 16, 2035	U-2471		
<u>LASMIDITAN SUCCINATE - REYVOW</u>						
N 211280	001	11053214	Dec 05, 2037	DS DP U-1719	NCE	Jan 31, 2025
		12071423	Jul 06, 2040	DP		
		7423050	Feb 17, 2028	DS DP U-1719		
<u>LASMIDITAN SUCCINATE - REYVOW</u>						
N 211280	002	11053214	Dec 05, 2037	DS DP U-1719	NCE	Jan 31, 2025
		12071423	Jul 06, 2040	DP		
		7423050	Feb 17, 2028	DS DP U-1719		
<u>LASMIDITAN SUCCINATE - REYVOW</u>						
N 211280	003	11053214	Dec 05, 2037	DS DP U-1719	NCE	Jan 31, 2025
		7423050	Feb 17, 2028	DS DP U-1719		
<u>LATANOPROST - XELPROS</u>						
N 206185	001	9539262	Oct 15, 2028	U-2400		
		9629852	Sep 12, 2029	DP		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>LATANOPROST - IYUZEH</u>						
N 216472	001 8637054	Jul 08, 2031	DP U-446			
<u>LATANOPROST; NETARSUDIL DIMESYLATE - ROCKLATAN</u>						
N 208259	001 10174017	Jan 27, 2030	DS DP U-1524			
	10532993	Jul 11, 2026	U-1524			
	10588901	Mar 14, 2034	DS DP U-1524			
	10654844	Jan 27, 2030	DS DP U-1524			
	10882840	Jul 11, 2026	U-1524			
	11020385	Mar 14, 2034	U-1524			
	11021456	Jul 11, 2026	U-1524			
	11028081	Jan 27, 2030	U-1524			
	11185538	Mar 14, 2034	DP			
	11197853	Mar 14, 2034	DP			
	11618748	Jan 27, 2030	U-1524			
	8394826	Nov 10, 2030	DS DP U-1524			
	8450344	Jul 11, 2026	DS DP U-1524			
	9096569	Jul 11, 2026	DS DP U-1524			
	9415043	Mar 14, 2034	DS			
	9931336	Mar 14, 2034	DS DP U-1524			
	9993470	Mar 14, 2034	DS DP U-1524			
<u>LATANOPROSTENE BUNOD - VYZULTA</u>						
N 207795	001 7273946	Oct 03, 2025	DS DP U-2144			
	7629345	Jan 05, 2025	DP U-2144			
	7910767	Jan 05, 2025	DS DP U-2144			
	8058467	Feb 21, 2029	DS U-2144			
<u>LAZERTINIB MESYLATE - LAZCLUZE</u>						
N 219008	001 11453656	Apr 18, 2038	DS DP		NCE	Aug 19, 2029
	11850248	Aug 01, 2041	U-3985			
	11879013	May 21, 2040	U-3985			
	11981659	Apr 18, 2038	U-3985			
	9593098	Oct 13, 2035	DS DP U-3985			
<u>LAZERTINIB MESYLATE - LAZCLUZE</u>						
N 219008	002 11453656	Apr 18, 2038	DS DP		NCE	Aug 19, 2029
	11850248	Aug 01, 2041	U-3985			
	11879013	May 21, 2040	U-3985			
	11981659	Apr 18, 2038	U-3985			
	9593098	Oct 13, 2035	DS DP U-3985			
<u>LEDIPASVIR; SOFOSBUVIR - HARVONI</u>						
N 205834	001 10039779	Jan 30, 2034	DS DP U-2369		ODE*	Aug 28, 2026
	10039779	Jan 30, 2034	DS DP U-2370			
	10039779*PED	Jul 30, 2034				
	10456414	Sep 14, 2032	DP			
	7964580	Mar 26, 2029	DS DP U-1470			
	7964580*PED	Sep 26, 2029				
	8088368	May 12, 2030	DS DP			
	8088368*PED	Nov 12, 2030				
	8273341	May 12, 2030	U-1470			
	8273341*PED	Nov 12, 2030				
	8334270	Mar 21, 2028	DS DP U-1470			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>LEDIPASVIR; SOFOSBUVIR - HARVONI</u>						
N 205834 001	8334270*PED	Sep 21, 2028				
	8580765	Mar 21, 2028	DS DP U-1470			
	8580765*PED	Sep 21, 2028				
	8618076	Dec 11, 2030	DS DP U-1470			
	8618076*PED	Jun 11, 2031				
	8633309	Mar 26, 2029	DS DP U-1470			
	8633309*PED	Sep 26, 2029				
	8735372	Mar 21, 2028	U-1470			
	8735372*PED	Sep 21, 2028				
	8822430	May 12, 2030	DS DP U-1470			
	8822430*PED	Nov 12, 2030				
	8841278	May 12, 2030	DP U-1470			
	8841278*PED	Nov 12, 2030				
	8889159	Mar 26, 2029	DP U-1470			
	8889159*PED	Sep 26, 2029				
	9085573	Mar 21, 2028	DS DP U-1470			
	9085573*PED	Sep 21, 2028				
	9284342	Sep 13, 2030	DS DP U-1470			
	9284342*PED	Mar 13, 2031				
	9393256	Sep 14, 2032	U-1470			
	9393256*PED	Mar 14, 2033				
	9511056	May 12, 2030	DP U-1470			
	9511056*PED	Nov 12, 2030				
<u>LEDIPASVIR; SOFOSBUVIR - HARVONI</u>						
N 205834 002	10039779	Jan 30, 2034	DS DP U-1470		ODE*	Aug 28, 2026
	10039779*PED	Jul 30, 2034				
	10456414	Sep 14, 2032	DP			
	7964580	Mar 26, 2029	DS DP U-1470			
	7964580*PED	Sep 26, 2029				
	8088368	May 12, 2030	DS DP			
	8088368*PED	Nov 12, 2030				
	8273341	May 12, 2030	U-1470			
	8273341*PED	Nov 12, 2030				
	8334270	Mar 21, 2028	DS DP U-1470			
	8334270*PED	Sep 21, 2028				
	8580765	Mar 21, 2028	DS DP U-1470			
	8580765*PED	Sep 21, 2028				
	8618076	Dec 11, 2030	DS DP U-1470			
	8618076*PED	Jun 11, 2031				
	8633309	Mar 26, 2029	DS DP U-1470			
	8633309*PED	Sep 26, 2029				
	8735372	Mar 21, 2028	U-1470			
	8735372*PED	Sep 21, 2028				
	8822430	May 12, 2030	DS DP U-1470			
	8822430*PED	Nov 12, 2030				
	8841278	May 12, 2030	DP U-1470			
	8841278*PED	Nov 12, 2030				
	8889159	Mar 26, 2029	DP U-1470			
	8889159*PED	Sep 26, 2029				

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>LEDIPASVIR; SOFOSBUVIR - HARVONI</u>						
N 205834	002 9085573	Mar 21, 2028	DS DP U-1470			
	9085573*PED	Sep 21, 2028				
	9284342	Sep 13, 2030	DS DP U-1470			
	9284342*PED	Mar 13, 2031				
	9393256	Sep 14, 2032	U-1470			
	9393256*PED	Mar 14, 2033				
	9511056	May 12, 2030	U-1470			
	9511056*PED	Nov 12, 2030				
<u>LEDIPASVIR; SOFOSBUVIR - HARVONI</u>						
N 212477	001 10456414	Sep 14, 2032	DP		ODE-262	Aug 28, 2026
	7964580	Mar 26, 2029	DS DP U-1470		ODE-263	Aug 28, 2026
	7964580*PED	Sep 26, 2029			ODE-264	Aug 28, 2026
	8088368	May 12, 2030	DS DP			
	8088368*PED	Nov 12, 2030				
	8273341	May 12, 2030	U-1470			
	8273341*PED	Nov 12, 2030				
	8334270	Mar 21, 2028	DS DP U-1470			
	8334270*PED	Sep 21, 2028				
	8580765	Mar 21, 2028	DS DP U-1470			
	8580765*PED	Sep 21, 2028				
	8618076	Dec 11, 2030	DS DP U-1470			
	8618076*PED	Jun 11, 2031				
	8633309	Mar 26, 2029	DS DP U-1470			
	8633309*PED	Sep 26, 2029				
	8735372	Mar 21, 2028	U-1470			
	8735372*PED	Sep 21, 2028				
	8822430	May 12, 2030	DS DP U-1470			
	8822430*PED	Nov 12, 2030				
	8841278	May 12, 2030	DP U-1470			
	8841278*PED	Nov 12, 2030				
	8889159	Mar 26, 2029	DP U-1470			
	8889159*PED	Sep 26, 2029				
	9085573	Mar 21, 2028	DS DP U-1470			
	9085573*PED	Sep 21, 2028				
	9284342	Sep 13, 2030	DS DP U-1470			
	9284342*PED	Mar 13, 2031				
	9393256	Sep 14, 2032	U-1470			
	9393256*PED	Mar 14, 2033				
	9511056	May 12, 2030	U-1470			
	9511056*PED	Nov 12, 2030				
<u>LEDIPASVIR; SOFOSBUVIR - HARVONI</u>						
N 212477	002 10456414	Sep 14, 2032	DP		ODE-262	Aug 28, 2026
	7964580	Mar 26, 2029	DS DP U-1470		ODE-263	Aug 28, 2026
	7964580*PED	Sep 26, 2029			ODE-264	Aug 28, 2026
	8088368	May 12, 2030	DS DP			
	8088368*PED	Nov 12, 2030				
	8273341	May 12, 2030	U-1470			
	8273341*PED	Nov 12, 2030				
	8334270	Mar 21, 2028	DS DP U-1470			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>LEDIPASVIR; SOFOSBUVIR - HARVONI</u>						
N 212477 002	8334270*PED	Sep 21, 2028				
	8580765	Mar 21, 2028	DS DP U-1470			
	8580765*PED	Sep 21, 2028				
	8618076	Dec 11, 2030	DS DP U-1470			
	8618076*PED	Jun 11, 2031				
	8633309	Mar 26, 2029	DS DP U-1470			
	8633309*PED	Sep 26, 2029				
	8735372	Mar 21, 2028	U-1470			
	8735372*PED	Sep 21, 2028				
	8822430	May 12, 2030	DS DP U-1470			
	8822430*PED	Nov 12, 2030				
	8841278	May 12, 2030	DP U-1470			
	8841278*PED	Nov 12, 2030				
	8889159	Mar 26, 2029	DP U-1470			
	8889159*PED	Sep 26, 2029				
	9085573	Mar 21, 2028	DS DP U-1470			
	9085573*PED	Sep 21, 2028				
	9284342	Sep 13, 2030	DS DP U-1470			
	9284342*PED	Mar 13, 2031				
	9393256	Sep 14, 2032	U-1470			
	9393256*PED	Mar 14, 2033				
	9511056	May 12, 2030	U-1470			
	9511056*PED	Nov 12, 2030				
<u>LEFAMULIN ACETATE - XENLETA</u>						
N 211672 001	8071643	Jan 16, 2029	DS DP		NCE	Aug 19, 2024
	8153689	Mar 19, 2028	DS DP		GAIN	Aug 19, 2029
	9120727	May 23, 2031	DS DP			
<u>LEFAMULIN ACETATE - XENLETA</u>						
N 211673 001	8071643	Jan 16, 2029	DS DP		NCE	Aug 19, 2024
	8153689	Mar 19, 2028	DS DP		GAIN	Aug 19, 2029
<u>LEMBorexant - DAYVIGO</u>						
N 212028 001	10188652	Oct 21, 2035	DP U-2791		M-293	Apr 20, 2026
	10702529	Oct 21, 2035	DP		NCE	Apr 07, 2025
	11026944	Oct 21, 2035	DP			
	8268848	Sep 20, 2031	DS DP U-2791			
<u>LEMBorexant - DAYVIGO</u>						
N 212028 002	10188652	Oct 21, 2035	DP U-2791		M-293	Apr 20, 2026
	10702529	Oct 21, 2035	DP		NCE	Apr 07, 2025
	11026944	Oct 21, 2035	DP			
	8268848	Sep 20, 2031	DS DP U-2791			
<u>LENACAPAVIR SODIUM - SUNLENCA</u>						
N 215973 001	10071985	Aug 17, 2037	DS DP		NCE	Dec 22, 2027
	10654827	Aug 17, 2037	U-3507			
	11267799	Aug 16, 2038	DS			
	11944611	Aug 28, 2041	U-3507			
	9951043	Feb 28, 2034	DS DP U-3507			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>LENACAPAVIR SODIUM - SUNLENCA</u>						
N 215974	001	10071985	Aug 17, 2037	DS DP	NCE	Dec 22, 2027
		10654827	Aug 17, 2037	U-3507		
		11267799	Aug 16, 2038	DS		
		11944611	Aug 28, 2041	U-3507		
		9951043	Feb 28, 2034	DS DP U-3507		
<u>LENALIDOMIDE - REVLIMID</u>						
N 021880	001	7465800	Apr 27, 2027	DS DP	ODE-241	May 28, 2026
		8741929	Mar 08, 2028	U-1983	ODE-245	May 28, 2026
<u>LENALIDOMIDE - REVLIMID</u>						
N 021880	002	7465800	Apr 27, 2027	DS DP	ODE-241	May 28, 2026
		8741929	Mar 08, 2028	U-1983	ODE-245	May 28, 2026
<u>LENALIDOMIDE - REVLIMID</u>						
N 021880	003	7465800	Apr 27, 2027	DS DP	ODE-241	May 28, 2026
		8741929	Mar 08, 2028	U-1983	ODE-245	May 28, 2026
<u>LENALIDOMIDE - REVLIMID</u>						
N 021880	004	7465800	Apr 27, 2027	DS DP	ODE-241	May 28, 2026
		8741929	Mar 08, 2028	U-1983	ODE-245	May 28, 2026
<u>LENALIDOMIDE - REVLIMID</u>						
N 021880	005	7465800	Apr 27, 2027	DS DP	ODE-241	May 28, 2026
		8741929	Mar 08, 2028	U-1983	ODE-245	May 28, 2026
<u>LENALIDOMIDE - REVLIMID</u>						
N 021880	006	7465800	Apr 27, 2027	DS DP	ODE-241	May 28, 2026
		8741929	Mar 08, 2028	U-1983	ODE-245	May 28, 2026
<u>LENIOLISIB PHOSPHATE - JOENJA</u>						
N 217759	001	8653092	Feb 19, 2032	DS DP	NCE	Mar 24, 2028
					ODE-430	Mar 24, 2030
<u>LENVATINIB MESYLATE - LENVIMA</u>						
N 206947	001	10259791	Aug 26, 2035	DS	M-14	Apr 03, 2027
		10259791*PED	Feb 26, 2036		ODE-196	Aug 15, 2025
		10407393	Aug 26, 2035	DS	PED	Feb 15, 2026
		10407393*PED	Feb 26, 2036		PED	Oct 03, 2027
		11090386	Feb 23, 2036	U-3519		
		11090386*PED	Aug 23, 2036			
		11186547	Aug 26, 2035	DS		
		11186547*PED	Feb 26, 2036			
		12083112	Mar 03, 2036	U-3996		
		12083112	Mar 03, 2036	U-3997		
		7253286	Oct 24, 2025	DS DP		
		7253286*PED	Apr 24, 2026			
		7612208	Sep 19, 2026	DS DP		
		7612208*PED	Mar 19, 2027			
		9006256	Jul 27, 2027	U-1695		
		9006256*PED	Jan 27, 2028			
<u>LENVATINIB MESYLATE - LENVIMA</u>						
N 206947	002	10259791	Aug 26, 2035	DS	M-14	Apr 03, 2027
		10259791*PED	Feb 26, 2036		ODE-196	Aug 15, 2025

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>LENVATINIB MESYLATE - LENVIMA</u>						
N 206947 002	10407393	Aug 26, 2035	DS		PED	Feb 15, 2026
	10407393*PED	Feb 26, 2036			PED	Oct 03, 2027
	11090386	Feb 23, 2036		U-3519		
	11090386*PED	Aug 23, 2036				
	11186547	Aug 26, 2035	DS			
	11186547*PED	Feb 26, 2036				
	12083112	Mar 03, 2036		U-3996		
	12083112	Mar 03, 2036		U-3997		
	7253286	Oct 24, 2025	DS DP			
	7253286*PED	Apr 24, 2026				
	7612208	Sep 19, 2026	DS DP			
	7612208*PED	Mar 19, 2027				
	9006256	Jul 27, 2027		U-1695		
	9006256*PED	Jan 27, 2028				
<u>LESINURAD - ZURAMPIC</u>						
N 207988 001	10183012	Nov 26, 2028		U-2311		
	8003681	Aug 25, 2025	DS			
	8084483	Aug 17, 2029		U-1801		
	8283369	Nov 26, 2028		U-1802		
	8283369	Nov 26, 2028		U-1804		
	8357713	Dec 22, 2029	DP	U-1801		
	8357713	Dec 22, 2029	DP	U-1802		
	8357713	Dec 22, 2029	DP	U-1803		
	8546436	Feb 29, 2032	DS DP			
	8546437	Apr 29, 2029		U-1803		
	9216179	Aug 01, 2031		U-1806		
	9956205	Dec 28, 2031		U-2311		
<u>LETERMOVIR - PREVYMIS</u>						
N 209939 001	RE46791	Jan 18, 2029	DS DP		D-189	Aug 02, 2026
					I-916	Jun 05, 2026
					NPP	Aug 30, 2027
					ODE-423	Jun 05, 2030
					ODE-495	Aug 30, 2031
					ODE-497	Aug 30, 2031
<u>LETERMOVIR - PREVYMIS</u>						
N 209939 002	RE46791	Jan 18, 2029	DS DP		D-189	Aug 02, 2026
					I-916	Jun 05, 2026
					NPP	Aug 30, 2027
					ODE-423	Jun 05, 2030
					ODE-495	Aug 30, 2031
					ODE-497	Aug 30, 2031
<u>LETERMOVIR - PREVYMIS</u>						
N 209940 001	10603384	Feb 28, 2033	DP		D-189	Aug 02, 2026
	RE46791	Jan 18, 2029	DS DP		I-916	Jun 05, 2026
					NPP	Aug 30, 2027
					ODE-423	Jun 05, 2030
					ODE-495	Aug 30, 2031
					ODE-497	Aug 30, 2031

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>LETERMOVIR - PREVYMIS</u>						
N 209940	001	10603384	Feb 28, 2033	DP	D-189	Aug 02, 2026
	RE46791	Jan 18, 2029	DS DP		I-916	Jun 05, 2026
					NPP	Aug 30, 2027
					ODE-423	Jun 05, 2030
					ODE-495	Aug 30, 2031
					ODE-497	Aug 30, 2031
<u>LETERMOVIR - PREVYMIS</u>						
N 209940	002	10603384	Feb 28, 2033	DP	D-189	Aug 02, 2026
	RE46791	Jan 18, 2029	DS DP		I-916	Jun 05, 2026
					NPP	Aug 30, 2027
					ODE-423	Jun 05, 2030
					ODE-495	Aug 30, 2031
					ODE-497	Aug 30, 2031
<u>LETERMOVIR - PREVYMIS</u>						
N 219104	001	RE46791	Jan 18, 2029	DS DP	NP	Aug 30, 2027
					ODE*	Jun 05, 2030
					ODE-495	Aug 30, 2031
					ODE-497	Aug 30, 2031
<u>LETERMOVIR - PREVYMIS</u>						
N 219104	002	RE46791	Jan 18, 2029	DS DP	NP	Aug 30, 2027
					ODE*	Jun 05, 2030
					ODE-495	Aug 30, 2031
					ODE-497	Aug 30, 2031
<u>LETROZOLE; RIBOCICLIB SUCCINATE - KISOALI FEMARA CO-PACK (COPACKAGED)</u>						
N 209935	001	10799506	Apr 14, 2036	DP	I-951	Sep 17, 2027
		12064434	Apr 14, 2036	DP		
		8324225	Jun 17, 2028	DS DP		
		8415355	Mar 13, 2031	DS DP		
		8685980	May 25, 2030	DS DP		
		8962630	Dec 09, 2029	U-2505		
		8962630	Dec 09, 2029	U-3264		
		8962630	Dec 09, 2029	U-3998		
		9193732	Nov 09, 2031	DS DP		
		9416136	Aug 20, 2029	U-2505		
		9416136	Aug 20, 2029	U-3264		
		9416136	Aug 20, 2029	U-3998		
		9868739	Nov 09, 2031	U-2505		
		9868739	Nov 09, 2031	U-3264		
		9868739	Nov 09, 2031	U-3998		
<u>LEUPROLIDE ACETATE - LUPRON DEPOT-PED KIT</u>						
N 020263	009	8921326	Feb 05, 2031	DP	NS	Apr 14, 2026
		9617303	Mar 22, 2028	U-3611		
<u>LEUPROLIDE ACETATE - LUPRON DEPOT</u>						
N 020517	003	8921326	Feb 05, 2031	DP	U-1666	
		9617303	Mar 22, 2028	U-4001		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>LEUPROLIDE ACETATE - ELIGARD KIT</u>						
N 021343	001	11771841	Dec 22, 2041	DS DP		
		11931559	Dec 22, 2041	DS		
<u>LEUPROLIDE ACETATE - ELIGARD KIT</u>						
N 021379	001	11771841	Dec 22, 2041	DS DP		
		11931559	Dec 22, 2041	DS		
<u>LEUPROLIDE ACETATE - ELIGARD KIT</u>						
N 021488	001	11771841	Dec 22, 2041	DS DP		
		11931559	Dec 22, 2041	DS		
<u>LEUPROLIDE ACETATE - ELIGARD KIT</u>						
N 021731	001	11771841	Dec 22, 2041	DS DP		
		11931559	Dec 22, 2041	DS		
<u>LEUPROLIDE ACETATE - FENSOLVI KIT</u>						
N 213150	001	11771841	Dec 22, 2041	DS DP		
		11931559	Dec 22, 2041	DS		
<u>LEUPROLIDE MESYLATE - CAMCEVI KIT</u>						
N 211488	001	10646572	Jan 16, 2027	DP		
		11717555	Jan 01, 2039	DP		
		12133878	Dec 18, 2037	DP		
		9572857	Jan 16, 2027	DP		
		9744207	Jan 16, 2027	DP		
<u>LEVACETYLLUCINE - AQNEURSA</u>						
N 219132	001				NCE	Sep 24, 2029
					ODE-498	Sep 24, 2031
<u>LEVETIRACETAM - KEPPRA</u>						
N 021035	001	8802142	Jun 07, 2031	DP		
		8802142*PED	Dec 07, 2031			
<u>LEVETIRACETAM - KEPPRA</u>						
N 021035	002	8802142	Jun 07, 2031	DP		
		8802142*PED	Dec 07, 2031			
<u>LEVETIRACETAM - KEPPRA</u>						
N 021035	003	8802142	Jun 07, 2031	DP		
		8802142*PED	Dec 07, 2031			
<u>LEVETIRACETAM - KEPPRA</u>						
N 021035	004	8802142	Jun 07, 2031	DP		
		8802142*PED	Dec 07, 2031			
<u>LEVETIRACETAM - KEPPRA XR</u>						
N 022285	001	7858122	Sep 17, 2028	DP		
<u>LEVETIRACETAM - KEPPRA XR</u>						
N 022285	002	7858122	Sep 17, 2028	DP		
<u>LEVETIRACETAM - ELEPSIA XR</u>						
N 204417	001	8163306	Sep 03, 2027	DP		
		8425938	Feb 22, 2026	DP		
		8431156	Oct 31, 2027	DP		
		8470367	Oct 31, 2027	DP		
		8535717	Feb 22, 2026	DP		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>LEVETIRACETAM - ELEPSIA XR</u>						
N 204417	001	8163306	Sep 03, 2027	DP		
		8425938	Feb 22, 2026	DP		
		8431156	Oct 31, 2027	DP		
		8470367	Oct 31, 2027	DP		
		8535717	Feb 22, 2026	DP		
<u>LEVETIRACETAM - ELEPSIA XR</u>						
N 204417	002	8163306	Sep 03, 2027	DP		
		8425938	Feb 22, 2026	DP		
		8431156	Oct 31, 2027	DP		
		8470367	Oct 31, 2027	DP		
		8535717	Feb 22, 2026	DP		
<u>LEVETIRACETAM - SPRITAM</u>						
N 207958	001	11160786	Mar 14, 2034	DP		
		9339489	Mar 14, 2034	DP	U-1850	
		9669009	Mar 14, 2034		U-1850	
		9669009	Mar 14, 2034		U-2021	
		9669009	Mar 14, 2034		U-2022	
<u>LEVETIRACETAM - SPRITAM</u>						
N 207958	002	11160786	Mar 14, 2034	DP		
		9339489	Mar 14, 2034	DP	U-1850	
		9669009	Mar 14, 2034		U-1850	
		9669009	Mar 14, 2034		U-2021	
		9669009	Mar 14, 2034		U-2022	
<u>LEVETIRACETAM - SPRITAM</u>						
N 207958	003	11160786	Mar 14, 2034	DP		
		9339489	Mar 14, 2034	DP	U-1850	
		9669009	Mar 14, 2034		U-1850	
		9669009	Mar 14, 2034		U-2021	
		9669009	Mar 14, 2034		U-2022	
<u>LEVETIRACETAM - SPRITAM</u>						
N 207958	004	11160786	Mar 14, 2034	DP		
		9339489	Mar 14, 2034	DP	U-1850	
		9669009	Mar 14, 2034		U-1850	
		9669009	Mar 14, 2034		U-2021	
		9669009	Mar 14, 2034		U-2022	
<u>LEVOCETIRIZINE DIHYDROCHLORIDE - XYZAL ALLERGY 24HR</u>						
N 209090	001	8633194	Oct 16, 2027	DP		
<u>LEVODOPA - INBRIJA</u>						
N 209184	001	8545878	Nov 16, 2032	DP		
		8685442	Nov 16, 2032	DP		
		8945612	Nov 16, 2032	DP		
		9393210	Nov 16, 2032	DP		
		RE43711	Feb 03, 2029		U-2484	
<u>LEVOKETOCONAZOLE - RECORLEV</u>						
N 214133	001	10098877	Jan 10, 2026	U-3283	ODE-385	Dec 30, 2028
		10517868	Jan 10, 2026	U-3283		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>LEVOKETOCONAZOLE - RECORLEV</u>						
N 214133	001	10835530	Jan 10, 2026	U-3283		
		11020393	Mar 02, 2040	U-3282		
		11278547	Mar 02, 2040	U-3282		
		11478471	Jan 10, 2026	U-3283		
		11903940	Mar 02, 2040	U-3821		
		9918984	Jan 10, 2026	U-3283		
<u>LEVOLEUCOVORIN - KHAPZORY</u>						
N 211226	001	11541012	Mar 25, 2039	DP		
<u>LEVOLEUCOVORIN - KHAPZORY</u>						
N 211226	002	11541012	Mar 25, 2039	DP		
<u>LEVOMILNACIPRAN HYDROCHLORIDE - FETZIMA</u>						
N 204168	001	8481598	Mar 02, 2031	U-839	M-304	Mar 24, 2026
		8865937	May 23, 2032	DS DP		
		RE43879	Jan 11, 2026	U-839		
<u>LEVOMILNACIPRAN HYDROCHLORIDE - FETZIMA</u>						
N 204168	002	8481598	Mar 02, 2031	U-839	M-304	Mar 24, 2026
		8865937	May 23, 2032	DS DP		
		RE43879	Jan 11, 2026	U-839		
<u>LEVOMILNACIPRAN HYDROCHLORIDE - FETZIMA</u>						
N 204168	003	8481598	Mar 02, 2031	U-839	M-304	Mar 24, 2026
		8865937	May 23, 2032	DS DP		
		RE43879	Jan 11, 2026	U-839		
<u>LEVOMILNACIPRAN HYDROCHLORIDE - FETZIMA</u>						
N 204168	004	8481598	Mar 02, 2031	U-839	M-304	Mar 24, 2026
		8865937	May 23, 2032	DS DP		
		RE43879	Jan 11, 2026	U-839		
<u>LEVONORGESTREL - MIRENA</u>						
N 021225	001	10561524	Sep 16, 2029	U-2948		
		11850182	Sep 14, 2029	DP U-3819		
<u>LEVONORGESTREL - SKYLA</u>						
N 203159	001	10561524	Sep 16, 2029	U-2948		
		11628088	Feb 07, 2027	DP		
		11850182	Sep 14, 2029	DP U-3819		
<u>LEVONORGESTREL - LILETTA</u>						
N 206229	001	10028858	Mar 22, 2034	DP U-2348	I-917	Jun 29, 2026
		11090186	Oct 24, 2033	U-2348		
		11571328	Sep 07, 2040	DP		
		12004992	Oct 06, 2033	DS		
<u>LEVONORGESTREL - KYLEENA</u>						
N 208224	001	10561524	Sep 16, 2029	U-2948		
		11628088	Feb 07, 2027	DP		
		11850182	Sep 14, 2029	DP U-3819		
<u>LEVOTHYROXINE SODIUM - LEVOLET</u>						
N 021137	003	10231931	Mar 23, 2038	DP		
		10406108	Mar 23, 2038	DP		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>LEVOTHYROXINE SODIUM - LEVOLET</u>						
N 021137 004	10231931	Mar 23, 2038	DP			
	10406108	Mar 23, 2038	DP			
<u>LEVOTHYROXINE SODIUM - LEVOLET</u>						
N 021137 005	10231931	Mar 23, 2038	DP			
	10406108	Mar 23, 2038	DP			
<u>LEVOTHYROXINE SODIUM - LEVOLET</u>						
N 021137 006	10231931	Mar 23, 2038	DP			
	10406108	Mar 23, 2038	DP			
<u>LEVOTHYROXINE SODIUM - LEVOLET</u>						
N 021137 007	10231931	Mar 23, 2038	DP			
	10406108	Mar 23, 2038	DP			
<u>LEVOTHYROXINE SODIUM - LEVOLET</u>						
N 021137 008	10231931	Mar 23, 2038	DP			
	10406108	Mar 23, 2038	DP			
<u>LEVOTHYROXINE SODIUM - LEVOLET</u>						
N 021137 009	10231931	Mar 23, 2038	DP			
	10406108	Mar 23, 2038	DP			
<u>LEVOTHYROXINE SODIUM - LEVOLET</u>						
N 021137 010	10231931	Mar 23, 2038	DP			
	10406108	Mar 23, 2038	DP			
<u>LEVOTHYROXINE SODIUM - LEVOLET</u>						
N 021137 011	10231931	Mar 23, 2038	DP			
	10406108	Mar 23, 2038	DP			
<u>LEVOTHYROXINE SODIUM - LEVOLET</u>						
N 021137 012	10231931	Mar 23, 2038	DP			
	10406108	Mar 23, 2038	DP			
<u>LEVOTHYROXINE SODIUM - LEVOTHYROXINE SODIUM</u>						
N 202231 001	9006289	Oct 03, 2032	DP			
	9168238	Aug 29, 2032	DP			
	9168239	Aug 29, 2032	DP			
<u>LEVOTHYROXINE SODIUM - LEVOTHYROXINE SODIUM</u>						
N 202231 002	9006289	Oct 03, 2032	DP			
	9168238	Aug 29, 2032	DP			
	9168239	Aug 29, 2032	DP			
<u>LEVOTHYROXINE SODIUM - LEVOTHYROXINE SODIUM</u>						
N 202231 003	9006289	Oct 03, 2032	DP			
	9168238	Aug 29, 2032	DP			
	9168239	Aug 29, 2032	DP			
<u>LEVOTHYROXINE SODIUM - TIROSINT-SOL</u>						
N 206977 001	10537538	Feb 28, 2037	DP			
	11096913	Feb 28, 2037	DP			
	11241382	Sep 17, 2039			U-3757	
	11241382	Sep 17, 2039			U-3758	

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>LEVOTHYROXINE SODIUM - TIROSINT-SOL</u>						
N 206977 002	10537538	Feb 28, 2037	DP			
	11096913	Feb 28, 2037	DP			
	11241382	Sep 17, 2039		U-3757		
	11241382	Sep 17, 2039		U-3758		
<u>LEVOTHYROXINE SODIUM - TIROSINT-SOL</u>						
N 206977 003	10537538	Feb 28, 2037	DP			
	11096913	Feb 28, 2037	DP			
	11241382	Sep 17, 2039		U-3757		
	11241382	Sep 17, 2039		U-3758		
<u>LEVOTHYROXINE SODIUM - TIROSINT-SOL</u>						
N 206977 004	10537538	Feb 28, 2037	DP			
	11096913	Feb 28, 2037	DP			
	11241382	Sep 17, 2039		U-3757		
	11241382	Sep 17, 2039		U-3758		
<u>LEVOTHYROXINE SODIUM - TIROSINT-SOL</u>						
N 206977 005	10537538	Feb 28, 2037	DP			
	11096913	Feb 28, 2037	DP			
	11241382	Sep 17, 2039		U-3757		
	11241382	Sep 17, 2039		U-3758		
<u>LEVOTHYROXINE SODIUM - TIROSINT-SOL</u>						
N 206977 006	10537538	Feb 28, 2037	DP			
	11096913	Feb 28, 2037	DP			
	11241382	Sep 17, 2039		U-3757		
	11241382	Sep 17, 2039		U-3758		
<u>LEVOTHYROXINE SODIUM - TIROSINT-SOL</u>						
N 206977 007	10537538	Feb 28, 2037	DP			
	11096913	Feb 28, 2037	DP			
	11241382	Sep 17, 2039		U-3757		
	11241382	Sep 17, 2039		U-3758		
<u>LEVOTHYROXINE SODIUM - TIROSINT-SOL</u>						
N 206977 008	10537538	Feb 28, 2037	DP			
	11096913	Feb 28, 2037	DP			
	11241382	Sep 17, 2039		U-3757		
	11241382	Sep 17, 2039		U-3758		
<u>LEVOTHYROXINE SODIUM - TIROSINT-SOL</u>						
N 206977 009	10537538	Feb 28, 2037	DP			
	11096913	Feb 28, 2037	DP			
	11241382	Sep 17, 2039		U-3757		
	11241382	Sep 17, 2039		U-3758		
<u>LEVOTHYROXINE SODIUM - TIROSINT-SOL</u>						
N 206977 010	10537538	Feb 28, 2037	DP			
	11096913	Feb 28, 2037	DP			
	11241382	Sep 17, 2039		U-3757		
	11241382	Sep 17, 2039		U-3758		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>LEVOTHYROXINE SODIUM - TIROSINT-SOL</u>						
N 206977	011	10537538	Feb 28, 2037	DP		
		11096913	Feb 28, 2037	DP		
		11241382	Sep 17, 2039		U-3757	
		11241382	Sep 17, 2039		U-3758	
<u>LEVOTHYROXINE SODIUM - TIROSINT-SOL</u>						
N 206977	012	10537538	Feb 28, 2037	DP		
		11096913	Feb 28, 2037	DP		
		11241382	Sep 17, 2039		U-3757	
		11241382	Sep 17, 2039		U-3758	
<u>LEVOTHYROXINE SODIUM - TIROSINT-SOL</u>						
N 206977	013	10537538	Feb 28, 2037	DP		
		11096913	Feb 28, 2037	DP		
		11241382	Sep 17, 2039		U-3757	
		11241382	Sep 17, 2039		U-3758	
<u>LEVOTHYROXINE SODIUM - TIROSINT-SOL</u>						
N 206977	014	10537538	Feb 28, 2037	DP		
		11096913	Feb 28, 2037	DP		
		11241382	Sep 17, 2039		U-3757	
		11241382	Sep 17, 2039		U-3758	
<u>LEVOTHYROXINE SODIUM - TIROSINT-SOL</u>						
N 206977	015	10537538	Feb 28, 2037	DP		
		11096913	Feb 28, 2037	DP		
		11241382	Sep 17, 2039		U-3757	
		11241382	Sep 17, 2039		U-3758	
<u>LEVOTHYROXINE SODIUM - LEVOTHYROXINE SODIUM</u>						
N 210632	001	10398669	Dec 01, 2036	DP		
		11135190	Dec 01, 2036	DP		
		9782376	Dec 01, 2036	DP		
<u>LEVOTHYROXINE SODIUM - LEVOTHYROXINE SODIUM</u>						
N 210632	002	10398669	Dec 01, 2036	DP		
		11135190	Dec 01, 2036	DP		
		9782376	Dec 01, 2036	DP		
<u>LEVOTHYROXINE SODIUM - LEVOTHYROXINE SODIUM</u>						
N 210632	003	10398669	Dec 01, 2036	DP		
		11135190	Dec 01, 2036	DP		
		9782376	Dec 01, 2036	DP		
<u>LEVOTHYROXINE SODIUM - THYQUIDITY</u>						
N 214047	001	9050307	Aug 06, 2031	DP		
<u>LEVOTHYROXINE SODIUM - LEVOTHYROXINE SODIUM</u>						
N 214253	001	11154498	Jul 20, 2036	DP		
<u>LEVOTHYROXINE SODIUM - ERMEZA</u>						
N 215809	001	9345772	Feb 27, 2035	DP		
<u>LIDOCAINE - ZTLIDO</u>						
N 207962	001	10765640	May 10, 2031	DP		
		10765749	May 10, 2031	DP		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>LIDOCAINE - ZTLIDO</u>						
N 207962	001 11278623	May 10, 2031	DP			
	11786455	May 10, 2031	DP	U-2267		
	11793766	May 10, 2031		U-2267		
	9283174	May 10, 2031	DP			
	9925264	May 10, 2031	DP	U-2267		
	9931403	May 10, 2031	DP			
<u>LIDOCAINE HYDROCHLORIDE - ZINGO</u>						
N 022114	001 8540665	Oct 22, 2029		U-1438		
	9358338	Apr 27, 2035		U-1870		
	9370622	Sep 28, 2035		U-1870		
<u>LIDOCAINE HYDROCHLORIDE - AKTEN</u>						
N 022221	001 8759401	Jul 24, 2026	DP	U-1523		
<u>LIDOCAINE; TETRACAINE - PLIAGLIS</u>						
N 021717	001 10350180	Jan 14, 2031	DP			
	10603293	Jan 14, 2031	DP			
	10751305	Jan 14, 2031	DP			
<u>LIFITEGRAST - XIIDRA</u>						
N 208073	001 11058677	Dec 18, 2033	DP			
	7314938	Mar 10, 2025	DS DP			
	8084047	May 17, 2026	DS DP			
	8168655	May 09, 2029		U-1880		
	8367701	Apr 15, 2029	DP	U-1880		
	8592450	May 17, 2026		U-1880		
	8927574	Nov 12, 2030	DP			
	9085553	Jul 25, 2033	DP			
	9353088	Oct 21, 2030	DP			
	9447077	Apr 15, 2029		U-1900		
	9890141	Oct 21, 2030	DS			
<u>LINACLOTIDE - LINZESS</u>						
N 202811	001 7304036	Aug 30, 2026	DS DP	U-1278	I-921	Jun 12, 2026
	7304036	Aug 30, 2026	DS DP	U-1516		
	8748573	Oct 30, 2031		U-1515		
	8748573	Oct 30, 2031		U-1516		
	8802628	Oct 30, 2031	DP			
	8933030	Feb 17, 2031	DP			
	9708371	Aug 16, 2033	DP	U-1515		
	9708371	Aug 16, 2033	DP	U-1516		
<u>LINACLOTIDE - LINZESS</u>						
N 202811	002 7304036	Aug 30, 2026	DS DP	U-1278		
	7304036	Aug 30, 2026	DS DP	U-1516		
	8748573	Oct 30, 2031		U-1515		
	8748573	Oct 30, 2031		U-1516		
	8802628	Oct 30, 2031	DP			
	8933030	Feb 17, 2031	DP			
	9708371	Aug 16, 2033	DP	U-1515		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>LINACLOTIDE - LINZESS</u>						
N 202811 003	10675325	Aug 11, 2031	DP		I-921	Jun 12, 2026
	10702576	Aug 11, 2031	U-1516			
	10702576	Aug 11, 2031	U-3644			
	7304036	Aug 30, 2026	DS DP U-1516			
	7304036	Aug 30, 2026	DS DP U-3644			
	8933030	Feb 17, 2031	DP U-1516			
	8933030	Feb 17, 2031	DP U-3644			
	9708371	Aug 16, 2033	DP U-1516			
	9708371	Aug 16, 2033	DP U-3644			
<u>LINAGLIPTIN - TRADJENTA</u>						
N 201280 001	10034877	Aug 05, 2029	U-2347		M-295	Jun 20, 2026
	10034877*PED	Feb 05, 2030			PED	Dec 20, 2026
	11033552	May 04, 2027	DP			
	11033552*PED	Nov 04, 2027				
	11911388	Apr 10, 2030	U-3854			
	7407955	May 02, 2025	DS DP			
	7407955*PED	Nov 02, 2025				
	8673927	May 04, 2027	U-1503	Y		
	8673927*PED	Nov 04, 2027				
	8846695	Jun 04, 2030	U-1503	Y		
	8846695*PED	Dec 04, 2030				
	8853156	Mar 05, 2031	DP U-1642			
	8853156*PED	Sep 05, 2031				
	8883805	Nov 26, 2025	DP			
	8883805*PED	May 26, 2026				
	9486526	Aug 05, 2029	U-1915			
	9486526*PED	Feb 05, 2030				
<u>LINAGLIPTIN; METFORMIN HYDROCHLORIDE - JENTADUETO</u>						
N 201281 001	10022379	Apr 02, 2029	U-2339		M-295	Jun 20, 2026
	10022379*PED	Oct 02, 2029			PED	Dec 20, 2026
	10973827	Apr 02, 2029	DP			
	10973827*PED	Oct 02, 2029				
	11911388	Apr 10, 2030	U-3854			
	7407955	May 02, 2025	DS DP			
	7407955*PED	Nov 02, 2025				
	8673927	May 04, 2027	U-1503	Y		
	8673927*PED	Nov 04, 2027				
	8846695	Jun 04, 2030	U-1503			
	8846695*PED	Dec 04, 2030				
	8883805	Nov 26, 2025	DP			
	8883805*PED	May 26, 2026				
	9155705	May 21, 2030	DP			
	9155705*PED	Nov 21, 2030				
	9415016	Apr 02, 2029	DP			
	9415016*PED	Oct 02, 2029				
<u>LINAGLIPTIN; METFORMIN HYDROCHLORIDE - JENTADUETO</u>						
N 201281 002	10022379	Apr 02, 2029	U-2339		M-295	Jun 20, 2026
	10022379*PED	Oct 02, 2029			PED	Dec 20, 2026

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>LINAGLIPTIN; METFORMIN HYDROCHLORIDE - JENTADUETO</u>						
N 201281 002	10973827	Apr 02, 2029	DP			
	10973827*PED	Oct 02, 2029				
	11911388	Apr 10, 2030		U-3854		
	7407955	May 02, 2025	DS DP			
	7407955*PED	Nov 02, 2025				
	8673927	May 04, 2027		U-1503	Y	
	8673927*PED	Nov 04, 2027				
	8846695	Jun 04, 2030		U-1503		
	8846695*PED	Dec 04, 2030				
	8883805	Nov 26, 2025	DP			
	8883805*PED	May 26, 2026				
	9155705	May 21, 2030	DP			
	9155705*PED	Nov 21, 2030				
	9415016	Apr 02, 2029	DP			
	9415016*PED	Oct 02, 2029				
<u>LINAGLIPTIN; METFORMIN HYDROCHLORIDE - JENTADUETO</u>						
N 201281 003	10022379	Apr 02, 2029		U-2339	M-295	Jun 20, 2026
	10022379*PED	Oct 02, 2029			PED	Dec 20, 2026
	10973827	Apr 02, 2029	DP			
	10973827*PED	Oct 02, 2029				
	11911388	Apr 10, 2030		U-3854		
	7407955	May 02, 2025	DS DP			
	7407955*PED	Nov 02, 2025				
	8673927	May 04, 2027		U-1503	Y	
	8673927*PED	Nov 04, 2027				
	8846695	Jun 04, 2030		U-1503	Y	
	8846695*PED	Dec 04, 2030				
	8883805	Nov 26, 2025	DP			
	8883805*PED	May 26, 2026				
	9155705	May 21, 2030	DP			
	9155705*PED	Nov 21, 2030				
	9415016	Apr 02, 2029	DP			
	9415016*PED	Oct 02, 2029				
<u>LINAGLIPTIN; METFORMIN HYDROCHLORIDE - JENTADUETO XR</u>						
N 208026 001	10022379	Apr 02, 2029		U-2339	M-295	Jun 20, 2026
	10022379*PED	Oct 02, 2029			PED	Dec 20, 2026
	11911388	Apr 10, 2030		U-3854		
	7407955	May 02, 2025	DS DP			
	7407955*PED	Nov 02, 2025				
	8673927	May 04, 2027		U-1503	Y	
	8673927*PED	Nov 04, 2027				
	8883805	Nov 26, 2025	DP			
	8883805*PED	May 26, 2026				
	9155705	May 21, 2030	DP			
	9155705*PED	Nov 21, 2030				
	9173859	May 04, 2027	DP U-1503		Y	
	9173859*PED	Nov 04, 2027				
	9415016	Apr 02, 2029	DP			
	9415016*PED	Oct 02, 2029				

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>LINAGLIPTIN; METFORMIN HYDROCHLORIDE - JENTADUETO XR</u>						
N 208026 001	9555001	Mar 06, 2033	DP U-1967			
	9555001	Mar 06, 2033	DP U-1968			
	9555001*PED	Sep 06, 2033				
<u>LINAGLIPTIN; METFORMIN HYDROCHLORIDE - JENTADUETO XR</u>						
N 208026 002	10022379	Apr 02, 2029	U-2339		M-295	Jun 20, 2026
	10022379*PED	Oct 02, 2029			PED	Dec 20, 2026
	11911388	Apr 10, 2030	U-3854			
	7407955	May 02, 2025	DS DP			
	7407955*PED	Nov 02, 2025				
	8673927	May 04, 2027	U-1503	Y		
	8673927*PED	Nov 04, 2027				
	8883805	Nov 26, 2025	DP			
	8883805*PED	May 26, 2026				
	9155705	May 21, 2030	DP			
	9155705*PED	Nov 21, 2030				
	9173859	May 04, 2027	DP U-1503	Y		
	9173859*PED	Nov 04, 2027				
	9415016	Apr 02, 2029	DP			
	9415016*PED	Oct 02, 2029				
	9555001	Mar 06, 2033	DP U-1967			
	9555001	Mar 06, 2033	DP U-1968			
	9555001*PED	Sep 06, 2033				
<u>LIRAGLUTIDE - VICTOZA</u>						
N 022341 001	8114833	Aug 13, 2025	DS DP			
	8114833*PED	Feb 13, 2026				
	9265893	Sep 23, 2032	DP			
	9265893*PED	Mar 23, 2033				
	9968659	Jan 09, 2037	U-2313			
	9968659*PED	Jul 09, 2037				
<u>LIRAGLUTIDE - SAXENDA</u>						
N 206321 001	10220155	Jul 17, 2026	DP			
	10220155*PED	Jan 17, 2027				
	10357616	Jan 20, 2026	DP			
	10376652	Jan 20, 2026	DP			
	11097063	Jul 17, 2026	DP			
	11311679	Jan 20, 2026	DP			
	11446443	Oct 20, 2025	DP			
	8114833	Aug 13, 2025	DP			
	8114833*PED	Feb 13, 2026				
	8684969	Oct 20, 2025	DP			
	8684969*PED	Apr 20, 2026				
	8920383	Jul 17, 2026	DP			
	8920383*PED	Jan 17, 2027				
	9108002	Jan 26, 2026	DP			
	9108002*PED	Jul 26, 2026				
	9132239	Feb 01, 2032	DP			
	9132239*PED	Aug 01, 2032				
	9457154	Sep 27, 2027	DP			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>LIRAGLUTIDE - SAXENDA</u>						
N 206321 001	9457154*PED	Mar 27, 2028				
	9616180	Jan 20, 2026	DP			
	9616180*PED	Jul 20, 2026				
	9687611	Feb 27, 2027	DP			
	9687611*PED	Aug 27, 2027				
	9775953	Jul 17, 2026	DP			
	9775953*PED	Jan 17, 2027				
	9861757	Jan 20, 2026	DP			
	9861757*PED	Jul 20, 2026				
	9968659	Jan 09, 2037		U-2438		
	9968659*PED	Jul 09, 2037				
	RE46363	Aug 03, 2026	DP			
	RE46363*PED	Feb 03, 2027				
<u>LISINAPRIL - OBRELIS</u>						
N 208401 001	10039800	Nov 06, 2035		U-1723		
	10039800	Nov 06, 2035		U-185		
	10039800	Nov 06, 2035		U-1864		
	10039800	Nov 06, 2035		U-1991		
	10039800	Nov 06, 2035		U-3		
	10039800	Nov 06, 2035		U-71		
	10039800	Nov 06, 2035		U-8		
	10265370	Nov 06, 2035	DP			
	10406199	Nov 06, 2035		U-1723		
	10406199	Nov 06, 2035		U-185		
	10406199	Nov 06, 2035		U-1864		
	10406199	Nov 06, 2035		U-1991		
	10406199	Nov 06, 2035		U-3		
	10406199	Nov 06, 2035		U-71		
	10406199	Nov 06, 2035		U-8		
	10940177	Nov 06, 2035	DP			
	11179434	Nov 06, 2035	DP			
	11771733	Nov 06, 2035	DP			
	12128083	Nov 06, 2035	DP			
	9463183	Nov 06, 2035	DP			
	9616096	Nov 06, 2035		U-1723		
	9616096	Nov 06, 2035		U-185		
	9616096	Nov 06, 2035		U-1864		
	9616096	Nov 06, 2035		U-1991		
	9616096	Nov 06, 2035		U-3		
	9616096	Nov 06, 2035		U-71		
	9616096	Nov 06, 2035		U-8		
	9814751	Nov 06, 2035	DP			
<u>LOFEXIDINE HYDROCHLORIDE - LOFEXIDINE HYDROCHLORIDE</u>						
A 218613 001					CGT	Feb 24, 2025
<u>LOMITAPIDE MESYLATE - JUXTAPID</u>						
N 203858 001	10016404	Mar 07, 2025		U-1316		
	10555938	Mar 07, 2025		U-1316		
	7932268	Aug 19, 2027		U-1316		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>LOMITAPIDE MESYLATE - JUXTAPID</u>						
N 203858	001	8618135	Mar 07, 2025	U-1316		
		9265758	Mar 07, 2025	U-1316		
		9364470	Mar 07, 2025	U-1851		
		9433617	Mar 07, 2025	U-1316		
		9861622	Mar 07, 2025	U-1316		
<u>LOMITAPIDE MESYLATE - JUXTAPID</u>						
N 203858	002	10016404	Mar 07, 2025	U-1316		
		10555938	Mar 07, 2025	U-1316		
		7932268	Aug 19, 2027	U-1316		
		8618135	Mar 07, 2025	U-1316		
		9265758	Mar 07, 2025	U-1316		
		9364470	Mar 07, 2025	U-1851		
		9433617	Mar 07, 2025	U-1316		
		9861622	Mar 07, 2025	U-1316		
<u>LOMITAPIDE MESYLATE - JUXTAPID</u>						
N 203858	003	10016404	Mar 07, 2025	U-1316		
		10555938	Mar 07, 2025	U-1316		
		7932268	Aug 19, 2027	U-1316		
		8618135	Mar 07, 2025	U-1316		
		9265758	Mar 07, 2025	U-1316		
		9364470	Mar 07, 2025	U-1851		
		9433617	Mar 07, 2025	U-1316		
		9861622	Mar 07, 2025	U-1316		
<u>LOMITAPIDE MESYLATE - JUXTAPID</u>						
N 203858	004	10016404	Mar 07, 2025	U-1316		
		10555938	Mar 07, 2025	U-1316		
		7932268	Aug 19, 2027	U-1316		
		8618135	Mar 07, 2025	U-1316		
		9265758	Mar 07, 2025	U-1316		
		9364470	Mar 07, 2025	U-1851		
		9433617	Mar 07, 2025	U-1316		
		9861622	Mar 07, 2025	U-1316		
<u>LOMITAPIDE MESYLATE - JUXTAPID</u>						
N 203858	005	10016404	Mar 07, 2025	U-1316		
		10555938	Mar 07, 2025	U-1316		
		7932268	Aug 19, 2027	U-1316		
		8618135	Mar 07, 2025	U-1316		
		9265758	Mar 07, 2025	U-1316		
		9364470	Mar 07, 2025	U-1851		
		9433617	Mar 07, 2025	U-1316		
		9861622	Mar 07, 2025	U-1316		
<u>LOMITAPIDE MESYLATE - JUXTAPID</u>						
N 203858	006	10016404	Mar 07, 2025	U-1316		
		10555938	Mar 07, 2025	U-1316		
		7932268	Aug 19, 2027	U-1316		
		8618135	Mar 07, 2025	U-1316		
		9265758	Mar 07, 2025	U-1316		
		9364470	Mar 07, 2025	U-1851		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>LOMITAPIDE MESYLATE - JUXTAPID</u>						
N 203858	006 9433617	Mar 07, 2025	U-1316			
	9861622	Mar 07, 2025	U-1316			
<u>LONAFARNIB - ZOKINVY</u>						
N 213969	001 7838531	Jul 26, 2025	U-3070		NCE	Nov 20, 2025
	8828356	Oct 17, 2025	U-3070		ODE-324	Nov 20, 2027
<u>LONAFARNIB - ZOKINVY</u>						
N 213969	002 7838531	Jul 26, 2025	U-3070		NCE	Nov 20, 2025
	8828356	Oct 17, 2025	U-3070		ODE-324	Nov 20, 2027
<u>LORAZEPAM - LOREEV XR</u>						
N 214826	001 8999393	Jan 08, 2034	DP U-3210			
<u>LORAZEPAM - LOREEV XR</u>						
N 214826	002 8999393	Jan 08, 2034	DP U-3210			
<u>LORAZEPAM - LOREEV XR</u>						
N 214826	003 8999393	Jan 08, 2034	DP U-3210			
<u>LORAZEPAM - LOREEV XR</u>						
N 214826	004 8999393	Jan 08, 2034	DP U-3210			
<u>LORLATINIB - LORBRENA</u>						
N 210868	001 10420749	Jul 27, 2036	DS DP U-2633		ODE-217	Nov 02, 2025
	10420749	Jul 27, 2036	DS DP U-3096		ODE-218	Nov 02, 2025
	11020376	Jul 27, 2036	DP		ODE-219	Nov 02, 2025
	11299500	Oct 04, 2038	DS		ODE-349	Mar 03, 2028
	8680111	Mar 05, 2033	DS DP			
<u>LORLATINIB - LORBRENA</u>						
N 210868	002 10420749	Jul 27, 2036	DS DP U-2633		ODE-217	Nov 02, 2025
	10420749	Jul 27, 2036	DS DP U-3096		ODE-218	Nov 02, 2025
	11020376	Jul 27, 2036	DP		ODE-219	Nov 02, 2025
	11299500	Oct 04, 2038	DS		ODE-349	Mar 03, 2028
	8680111	Mar 05, 2033	DS DP			
<u>LOTEPREDNOL ETABONATE - LOTEMAX SM</u>						
N 208219	001 10596107	Dec 23, 2036	DP U-2764			
	11534395	Jan 26, 2036	DP U-2764			
<u>LOTEPREDNOL ETABONATE - INVELTYS</u>						
N 210565	001 10058511	May 03, 2033	DP U-2492			
	10646437	May 03, 2033	DP			
	10688045	May 03, 2033	DP			
	10864219	May 03, 2033	U-3011			
	11219597	May 03, 2033	U-3278			
	11219597	May 03, 2033	U-3279			
	11642317	May 03, 2033	DP			
	11872318	May 03, 2033	DP			
	12115246	May 03, 2033	DP U-4025			
	9056057	May 03, 2033	DP U-2491			
	9393213	May 03, 2033	DP			
	9532955	May 03, 2033	U-2491			
	9737491	May 03, 2033	U-2492			
	9827191	May 03, 2033	DP U-2493			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>LOTEPREDNOL ETABONATE - INVELTYS</u>						
N 210565	001	10058511	May 03, 2033	DP	U-2492	
		10646437	May 03, 2033	DP		
		10688045	May 03, 2033	DP		
		10864219	May 03, 2033		U-3011	
		11219597	May 03, 2033		U-3278	
		11219597	May 03, 2033		U-3279	
		11642317	May 03, 2033	DP		
		11872318	May 03, 2033	DP		
		12115246	May 03, 2033	DP	U-4025	
		9056057	May 03, 2033	DP	U-2491	
		9393213	May 03, 2033	DP		
		9532955	May 03, 2033		U-2491	
		9737491	May 03, 2033		U-2492	
		9827191	May 03, 2033	DP	U-2493	
<u>LOTEPREDNOL ETABONATE - EYSUVIS</u>						
N 210933	001	10058511	May 03, 2033	DP	U-2492	
		10646436	May 03, 2033	DP		
		10688045	May 03, 2033	DP		
		10857096	May 03, 2033		U-2985	
		10940108	May 03, 2033		U-2985	
		10945948	May 03, 2033		U-2985	
		10993908	May 03, 2033		U-3117	
		11219596	May 03, 2033		U-2985	
		11596599	May 03, 2033		U-2985	
		11642317	May 03, 2033	DP		
		11872318	May 03, 2033	DP		
		12115246	May 03, 2033	DP	U-2985	
		9056057	May 03, 2033	DP	U-2491	
		9393213	May 03, 2033	DP		
		9532955	May 03, 2033		U-2491	
		9737491	May 03, 2033		U-2492	
		9827191	May 03, 2033	DP	U-2985	
<u>LOTILANER - XDEMVY</u>						
N 217603	001	10835517	Dec 14, 2038		U-3674	NCE Jul 24, 2028
		11197847	Dec 14, 2038		U-3674	
		11690826	Dec 14, 2038		U-3674	
		11690827	Dec 14, 2038		U-3674	
		11752137	Dec 14, 2038	DP		
		8383659	Jan 17, 2030	DS	DP	
<u>LOXAPINE - ADASUVE</u>						
N 022549	001	8387612	Oct 23, 2026	DP		
<u>LUBIPROSTONE - AMITIZA</u>						
N 021908	001	8026393	Oct 25, 2027	DP		
		8338639	Jan 23, 2027	DP		
		8748481	Sep 01, 2025		U-1520	
		8779187	Jan 23, 2027	DP		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>LUBIPROSTONE - AMITIZA</u>						
N 021908	002 8026393	Oct 25, 2027	DP			
	8338639	Jan 23, 2027	DP			
	8748481	Sep 01, 2025		U-1519		
	8779187	Jan 23, 2027	DP			
<u>LULICONAZOLE - LUZU</u>						
N 204153	001 8980931	Apr 28, 2034	DP			
	9012484	Sep 06, 2033	DS DP	U-540		
	9199977	Sep 06, 2033	DS DP			
	9453006	Sep 06, 2033	DS			
<u>LUMASIRAN SODIUM - OXLUMO</u>						
N 214103	001 10131907	Aug 24, 2028	DS DP	U-2995	I-901	Oct 06, 2025
	10435692	Dec 26, 2034		U-2995	NCE	Nov 23, 2025
	10465195	Dec 26, 2034	DS DP	U-2995	ODE-339	Nov 23, 2027
	10478500	Oct 09, 2035	DS DP	U-2995	ODE-415	Oct 06, 2029
	10487330	Dec 26, 2034	DS DP	U-2995		
	10612024	Aug 14, 2035	DS DP	U-2995		
	10612027	Aug 14, 2035	DS DP	U-2995		
	11060093	Dec 26, 2034	DS DP	U-2995		
	11261447	Nov 20, 2038	DS DP	U-2995		
	11401517	Aug 14, 2035	DS DP	U-2995		
	11446380	Oct 09, 2035	DS DP			
	8106022	Dec 12, 2029	DS DP	U-2995		
	8828956	Dec 04, 2028	DS DP	U-2995		
	9828606	Dec 26, 2034	DS DP			
<u>LUMATEPERONE TOSYLATE - CAPLYTA</u>						
N 209500	001 10117867	May 27, 2029		U-3271		
	10464938	Mar 12, 2028	DP			
	10695345	Aug 30, 2039	DP	U-543		
	10960009	Dec 03, 2034		U-814		
	11026951	Dec 03, 2034		U-3274		
	11690842	Aug 30, 2039	DP	U-3362		
	11690842	Aug 30, 2039	DP	U-3363		
	11753419	Dec 10, 2040	DP			
	11806348	Aug 30, 2039	DP	U-3362		
	11806348	Aug 30, 2039	DP	U-3363		
	11980617	Oct 27, 2039		U-3940		
	12070459	Aug 30, 2039	DP	U-3362		
	12070459	Aug 30, 2039	DP	U-3363		
	12090155	Jul 07, 2040	DP	U-4000		
	12122792	Dec 10, 2040	DP	U-4019		
	12122792	Dec 10, 2040	DP	U-4020		
	12128043	Aug 30, 2039	DP	U-3362		
	12128043	Aug 30, 2039	DP	U-3363		
	8598119	Dec 28, 2029		U-543	Y	
	8648077	Dec 01, 2029	DS DP			
	9199995	Mar 12, 2029		U-2713		
	9586960	Mar 12, 2029	DS DP			
	9616061	May 27, 2029	DP			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>LUMATEPERONE TOSYLATE - CAPLYTA</u>						
N 209500 001	9956227	Dec 03, 2034	U-2714			
	RE48825	Mar 12, 2029	DS DP			
	RE48839	Aug 19, 2033	U-3271			
	RE48839	Aug 19, 2033	U-814			
<u>LUMATEPERONE TOSYLATE - CAPLYTA</u>						
N 209500 002	10117867	May 27, 2029	U-3271			
	10464938	Mar 12, 2028	DP			
	10695345	Aug 30, 2039	DP U-814			
	11026951	Dec 03, 2034	U-3364			
	11052084	Aug 30, 2039	DP U-3362			
	11052084	Aug 30, 2039	DP U-3363			
	11690842	Aug 30, 2039	DP U-3362			
	11690842	Aug 30, 2039	DP U-3363			
	11753419	Dec 10, 2040	DP			
	11806348	Aug 30, 2039	DP U-3362			
	11806348	Aug 30, 2039	DP U-3363			
	11980617	Oct 27, 2039	U-3940			
	12122792	Dec 10, 2040	DP U-4019			
	12122792	Dec 10, 2040	DP U-4020			
	8648077	Dec 01, 2029	DS DP			
	9168258	May 27, 2029	DP			
	9199995	Mar 12, 2029	U-2713			
	9616061	May 27, 2029	DP			
	9956227	Dec 03, 2034	U-2714			
	RE48825	Feb 12, 2029	DS			
	RE48839	Aug 19, 2033	U-3271			
	RE48839	Aug 19, 2033	U-814			
<u>LUMATEPERONE TOSYLATE - CAPLYTA</u>						
N 209500 003	10117867	May 27, 2029	U-3271			
	10464938	Mar 12, 2028	DP			
	10695345	Aug 30, 2039	DP U-814			
	11026951	Dec 03, 2034	U-3364			
	11052084	Aug 30, 2039	DP U-3362			
	11052084	Aug 30, 2039	DP U-3363			
	11690842	Aug 30, 2039	DP U-3362			
	11690842	Aug 30, 2039	DP U-3363			
	11753419	Dec 10, 2040	DP			
	11806348	Aug 30, 2039	DP U-3362			
	11806348	Aug 30, 2039	DP U-3363			
	11980617	Oct 27, 2039	U-3940			
	12122792	Dec 10, 2040	DP U-4019			
	12122792	Dec 10, 2040	DP U-4020			
	8648077	Dec 01, 2029	DS DP			
	9199995	Mar 12, 2029	U-2713			
	9616061	May 27, 2029	DP			
	9956227	Dec 03, 2034	U-2714			
	RE48825	Feb 12, 2029	DS			
	RE48839	Aug 19, 2033	U-3271			
	RE48839	Aug 19, 2033	U-814			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>LUMATEPERONE TOSYLATE - CAPLYTA</u>						
N 209500 003	10117867	May 27, 2029	U-3271			
	10464938	Mar 12, 2028	DP			
	10695345	Aug 30, 2039	DP U-814			
	11026951	Dec 03, 2034	U-3364			
	11052084	Aug 30, 2039	DP U-3362			
	11052084	Aug 30, 2039	DP U-3363			
	11690842	Aug 30, 2039	DP U-3362			
	11690842	Aug 30, 2039	DP U-3363			
	11753419	Dec 10, 2040	DP			
	11806348	Aug 30, 2039	DP U-3362			
	11806348	Aug 30, 2039	DP U-3363			
	11980617	Oct 27, 2039	U-3940			
	12122792	Dec 10, 2040	DP U-4019			
	12122792	Dec 10, 2040	DP U-4020			
	8648077	Dec 01, 2029	DS DP			
	9199995	Mar 12, 2029	U-2713			
	9616061	May 27, 2029	DP			
	9956227	Dec 03, 2034	U-2714			
	RE48825	Feb 12, 2029	DS			
	RE48839	Aug 19, 2033	U-3271			
	RE48839	Aug 19, 2033	U-814			
<u>LURASIDONE HYDROCHLORIDE - LATUDA</u>						
N 200603 001	8729085	May 26, 2026	DP			
	8729085*PED	Nov 26, 2026				
	8883794	May 26, 2026	DP			
	8883794*PED	Nov 26, 2026				
	9259423	May 23, 2031	U-1822			
	9259423*PED	Nov 23, 2031				
	9555027	May 26, 2026	DP U-543			
	9827242	May 23, 2031	U-2199			
	9827242	May 23, 2031	U-2201			
	9907794	May 26, 2026	DP			
	9907794*PED	Nov 26, 2026				
	RE45573	Jun 23, 2025	DS			
	RE45573*PED	Dec 23, 2025				
<u>LURASIDONE HYDROCHLORIDE - LATUDA</u>						
N 200603 002	8729085	May 26, 2026	DP			
	8729085*PED	Nov 26, 2026				
	8883794	May 26, 2026	DP			
	8883794*PED	Nov 26, 2026				
	9259423	May 23, 2031	U-1822			
	9259423*PED	Nov 23, 2031				
	9555027	May 26, 2026	DP U-543			
	9827242	May 23, 2031	U-2199			
	9827242	May 23, 2031	U-2201			
	9907794	May 26, 2026	DP			
	9907794*PED	Nov 26, 2026				
	RE45573	Jun 23, 2025	DS			
	RE45573*PED	Dec 23, 2025				

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>LURASIDONE HYDROCHLORIDE - LATUDA</u>						
N 200603 002	8729085	May 26, 2026	DP			
	8729085*PED	Nov 26, 2026				
	8883794	May 26, 2026	DP			
	8883794*PED	Nov 26, 2026				
	9259423	May 23, 2031		U-1822		
	9259423*PED	Nov 23, 2031				
	9555027	May 26, 2026	DP	U-543		
	9827242	May 23, 2031		U-2199		
	9827242	May 23, 2031		U-2201		
	9907794	May 26, 2026	DP			
	9907794*PED	Nov 26, 2026				
	RE45573	Jun 23, 2025	DS			
	RE45573*PED	Dec 23, 2025				
<u>LURASIDONE HYDROCHLORIDE - LATUDA</u>						
N 200603 003	8729085	May 26, 2026	DP			
	8729085*PED	Nov 26, 2026				
	8883794	May 26, 2026	DP			
	8883794*PED	Nov 26, 2026				
	9259423	May 23, 2031		U-1822		
	9259423*PED	Nov 23, 2031				
	9555027	May 26, 2026	DP	U-543		
	9827242	May 23, 2031		U-2199		
	9827242	May 23, 2031		U-2201		
	9907794	May 26, 2026	DP			
	9907794*PED	Nov 26, 2026				
	RE45573	Jun 23, 2025	DS			
	RE45573*PED	Dec 23, 2025				
<u>LURASIDONE HYDROCHLORIDE - LATUDA</u>						
N 200603 004	8729085	May 26, 2026	DP			
	8729085*PED	Nov 26, 2026				
	8883794	May 26, 2026	DP			
	8883794*PED	Nov 26, 2026				
	9259423	May 23, 2031		U-1822		
	9259423*PED	Nov 23, 2031				
	9555027	May 26, 2026	DP	U-543		
	9827242	May 23, 2031		U-2199		
	9827242	May 23, 2031		U-2201		
	9907794	May 26, 2026	DP			
	9907794*PED	Nov 26, 2026				
	RE45573	Jun 23, 2025	DS			
	RE45573*PED	Dec 23, 2025				
<u>LURASIDONE HYDROCHLORIDE - LATUDA</u>						
N 200603 005	8729085	May 26, 2026	DP			
	8729085*PED	Nov 26, 2026				
	8883794	May 26, 2026	DP			
	8883794*PED	Nov 26, 2026				
	9259423	May 23, 2031		U-1822		
	9259423*PED	Nov 23, 2031				

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>LURASIDONE HYDROCHLORIDE - LATUDA</u>						
N 200603	005 9555027	May 26, 2026	DP U-543			
	9827242	May 23, 2031	U-2199			
	9827242	May 23, 2031	U-2201			
	9907794	May 26, 2026	DP			
	9907794*PED	Nov 26, 2026				
	RE45573	Jun 23, 2025	DS			
	RE45573*PED	Dec 23, 2025				
<u>LURBINECTEDIN - ZEPZELCA</u>						
N 213702	001 7763615	Dec 13, 2029	DS DP U-2836		NCE ODE-304	Jun 15, 2025 Jun 15, 2027
<u>LUSUTROMBOPAG - MULPLETA</u>						
N 210923	001 8530668	Jan 21, 2030	DS DP			
	8889722	Jul 29, 2028	DS DP			
	9427402	Sep 29, 2031	DP			
<u>LUTETIUM LU 177 DOTATATE - LUTATHERA</u>						
N 208700	001 10596276	Jul 25, 2038	DP		NPP	Apr 23, 2027
	10596276*PED	Jan 25, 2039			ODE-166	Jan 26, 2025
	10596278	Jul 25, 2038	DP		ODE-479	Apr 23, 2031
	10596278*PED	Jan 25, 2039			PED	Jul 26, 2025
	11904027	Jul 25, 2038	DP		PED	Oct 23, 2027
	11904027*PED	Jan 25, 2039			PED	Oct 23, 2031
	12144873	Jul 25, 2038	U-4036			
	12144873*PED	Jan 25, 2039				
	12151003	Jul 25, 2038	DP U-4036			
	12151003*PED	Jan 25, 2039				
	12161732	Jul 25, 2038	DP U-4036			
	12161732*PED	Jan 25, 2039				
	12168063	Jul 25, 2038	DP U-4036			
<u>LUTETIUM LU-177 VIPIVOTIDE TETRAJETAN - PLUVICTO</u>						
N 215833	001 10398791	Oct 17, 2034	DS DP		NCE	Mar 23, 2027
	10406240	Aug 15, 2028	DS DP U-3345			
	11318121	Aug 15, 2028	DS DP U-3345			
	11951190	Nov 12, 2035	U-3345			
<u>MACIMORELIN ACETATE - MACRILEN</u>						
N 205598	001 8192719	Oct 12, 2027	U-2220			
<u>MACITENTAN - OPSUMIT</u>						
N 204410	001 10946015	Sep 11, 2026	DP U-1445			
	7094781	Dec 05, 2025	DS DP			
	8268847	Apr 18, 2029	U-1446			
	8367685	Oct 04, 2028	DP U-1445			
	9265762	May 29, 2027	DP U-1820			
<u>MACITENTAN; TADALAFIL - OPSYNOVI</u>						
N 218490	001 10946015	Sep 11, 2026	DP U-3881		NP	Mar 22, 2027
	7094781	Dec 05, 2025	DS DP		ODE-475	Mar 22, 2031
	8268847	Apr 18, 2029	DP U-3882			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>MACITENTAN; TADALAFIL - OPSYNVI</u>						
N 218490	002 10946015	Sep 11, 2026	DP U-3881		NP	Mar 22, 2027
	7094781	Dec 05, 2025	DS DP		ODE-475	Mar 22, 2031
	8268847	Apr 18, 2029	DP U-3882			
<u>MAGNESIUM SULFATE; POLYETHYLENE GLYCOL 3350; POTASSIUM CHLORIDE; SODIUM CHLORIDE; SODIUM SULFATE - SUFLAVE</u>						
N 215344	001				NP	Jun 15, 2026
<u>MAGNESIUM SULFATE; POTASSIUM CHLORIDE; SODIUM SULFATE - SUTAB</u>						
N 213135	001 10143656	Aug 04, 2037	DP			
	11033498	Aug 04, 2037	U-3164			
	11382864	Aug 04, 2037	U-3164			
	11638697	Aug 04, 2037	DP			
<u>MAGNESIUM SULFATE; POTASSIUM SULFATE; SODIUM SULFATE - SUPREP BOWEL PREP KIT</u>						
N 022372	001				ODE-315	Aug 05, 2027
<u>MALATHION - OVIDE</u>						
N 018613	001 7560445	Feb 01, 2027	DS DP U-986			
	7977324	Aug 14, 2026	DP			
<u>MANNITOL - BRONCHITOL</u>						
N 202049	001				ODE-327	Oct 30, 2027
<u>MARALIXIBAT CHLORIDE - LIVMARLI</u>						
N 214662	001 10512657	Oct 26, 2032	U-3850		I-938	Mar 13, 2027
	11229647	Feb 12, 2040	U-3290		NCE	Sep 29, 2026
	11229647	Feb 12, 2040	U-3974		NPP	Mar 13, 2026
	11229661	Oct 26, 2032	U-3850		ODE-379	Sep 29, 2028
	11260053	May 26, 2031	U-3290		ODE-429	Mar 13, 2030
	11260053	May 26, 2031	U-3974		ODE-471	Mar 13, 2031
	11376251	Oct 26, 2032	U-3290			
	11376251	Oct 26, 2032	U-3974			
	11497745	Feb 12, 2040	U-3290			
	11497745	Feb 12, 2040	U-3974			
	11918578	Feb 12, 2040	U-3290			
	11918578	Feb 12, 2040	U-3974			
<u>MARALIXIBAT CHLORIDE - LIVMARLI</u>						
N 214662	002 10512657	Oct 26, 2032	U-3973		NCE	Sep 29, 2026
	11229661	Oct 26, 2032	U-3973		ODE*	Sep 29, 2028
					ODE*	Mar 13, 2030
					ODE*	Mar 13, 2031
					ODE-490	Jul 24, 2031
<u>MARIBAVIR - LIVTENCITY</u>						
N 215596	001 11684632	Jan 04, 2032	U-3650		NCE	Nov 23, 2026
					ODE-388	Nov 23, 2028
<u>MAVACAMTEN - CAMZYOS</u>						
N 214998	001 9585883	Jun 19, 2034	U-3373		M-297	Jun 15, 2026
	RE50050	Jun 19, 2034	DS DP		NCE	Apr 28, 2027
					ODE-398	Apr 28, 2029
<u>MAVACAMTEN - CAMZYOS</u>						
N 214998	002 9585883	Jun 19, 2034	U-3373		M-297	Jun 15, 2026
	RE50050	Jun 19, 2034	DS DP		NCE	Apr 28, 2027

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>MAVACAMTEN - CAMZYOS</u>						
N 214998	002				ODE-398	Apr 28, 2029
<u>MAVACAMTEN - CAMZYOS</u>						
N 214998	003	9585883	Jun 19, 2034	U-3373	M-297	Jun 15, 2026
	RE50050		Jun 19, 2034	DS DP	NCE	Apr 28, 2027
					ODE-398	Apr 28, 2029
<u>MAVACAMTEN - CAMZYOS</u>						
N 214998	004	9585883	Jun 19, 2034	U-3373	M-297	Jun 15, 2026
	RE50050		Jun 19, 2034	DS DP	NCE	Apr 28, 2027
					ODE-398	Apr 28, 2029
<u>MAVORIXAFOR - XOLREMDI</u>						
N 218709	001	10548889	Dec 11, 2038	DP	NCE	Apr 26, 2029
		10610527	Dec 22, 2036	U-3932	ODE-480	Apr 26, 2031
		10953003	Dec 14, 2036	DP		
		11045461	Dec 11, 2038	DP		
		11219621	Dec 22, 2036	U-3932		
		12115156	Dec 11, 2038	DP		
<u>MECHLORETHAMINE HYDROCHLORIDE - VALCHLOR</u>						
N 202317	001	7838564	Mar 07, 2026	DP		
		7872050	Jul 08, 2029	U-1427		
		8450375	Mar 07, 2026	DP		
		8501818	Mar 07, 2026	DP		
		8501819	Mar 07, 2026	U-1427		
		9382191	Mar 07, 2026	DP		
<u>MELOXICAM - VIVLODEX</u>						
N 207233	001	9526734	Mar 31, 2033	DP		
		9649318	Mar 31, 2035	DP		
		9808468	Mar 31, 2035	U-2160		
		9808468	Mar 31, 2035	U-2165		
<u>MELOXICAM - VIVLODEX</u>						
N 207233	002	9526734	Mar 31, 2033	DP		
		9649318	Mar 31, 2035	DP		
		9808468	Mar 31, 2035	U-2160		
		9808468	Mar 31, 2035	U-2165		
<u>MELOXICAM - ANJESO</u>						
N 210583	001	10709713	May 26, 2030	U-2750		
		10881663	Mar 08, 2039	U-3038		
		11253478	May 26, 2030	DP U-3318		
		11458145	Mar 08, 2039	U-3318		
		9974746	May 26, 2030	DP		
<u>MELOXICAM - OMIIZ ODT</u>						
N 211210	001	8545879	Aug 31, 2030	DP		
<u>MELOXICAM - OMIIZ ODT</u>						
N 211210	002	8545879	Aug 31, 2030	DP		
<u>MELPHALAN HYDROCHLORIDE - HEPZATO</u>						
N 201848	001	10098997	Nov 07, 2032	DP	NP	Aug 14, 2026
		10195334	Jan 16, 2033	DP	ODE-438	Aug 14, 2030

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>MELPHALAN HYDROCHLORIDE - HEPZATO</u>						
N 201848	001	10369264	Nov 07, 2032	DP		
		10569004	Nov 07, 2032		U-3680	
		10569004	Nov 07, 2032		U-3683	
		11083831	Dec 30, 2032	DP		
		11241522	Nov 07, 2032	DP		
		11633528	Nov 07, 2032		U-3675	
		11833286	Dec 30, 2032	DP		
		9314561	Feb 07, 2034	DP		
		9707331	Sep 17, 2034	DP		
<u>MELPHALAN HYDROCHLORIDE - EVOMELA</u>						
N 207155	001	10040872	Jan 30, 2034	DP		
		10864183	May 28, 2030	DP		
		10940128	Jun 14, 2030	DP	U-3086	
		11020363	May 28, 2030	DP		
		8410077	Mar 13, 2029	DP		
		9200088	Mar 13, 2029	DP		
		9493582	Feb 27, 2033	DP		
<u>MELPHALAN HYDROCHLORIDE - IVRA</u>						
N 217110	001	10537520	Jun 29, 2036	DP		
<u>MEMANTINE HYDROCHLORIDE - NAMENDA XR</u>						
N 022525	001	8039009	Mar 24, 2029		U-539	
		8039009*PED	Sep 24, 2029			
<u>MEMANTINE HYDROCHLORIDE - NAMENDA XR</u>						
N 022525	002	8039009	Mar 24, 2029		U-539	
		8039009*PED	Sep 24, 2029			
<u>MEMANTINE HYDROCHLORIDE - NAMENDA XR</u>						
N 022525	003	8039009	Mar 24, 2029		U-539	
		8039009*PED	Sep 24, 2029			
<u>MEMANTINE HYDROCHLORIDE - NAMENDA XR</u>						
N 022525	004	8039009	Mar 24, 2029		U-539	
		8039009*PED	Sep 24, 2029			
<u>MENTHOL; METHYL SALICYLATE - SALONPAS</u>						
N 022029	001	8809615	Jan 03, 2030	DP		
		9233184	Aug 01, 2027	DP		
<u>MENTHOL; METHYL SALICYLATE - SALONPAS</u>						
N 022029	002	8809615	Jan 03, 2030	DP		
		9233184	Aug 01, 2027	DP		
<u>MEROPENEM; VABORBACTAM - VABOMERE</u>						
N 209776	001	10172874	Aug 08, 2031	DP		NCE Aug 29, 2022
		10183034	Aug 08, 2031		U-2490	GAIN Aug 29, 2027
		10561675	Aug 08, 2031		U-2490	
		11007206	Aug 08, 2031		U-3128	
		11376237	Apr 06, 2039		U-3421	
		12171772	Aug 08, 2031	DS		
		8680136	Aug 29, 2031	DS DP		
		9694025	Aug 08, 2031		U-2120	

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>MESALAMINE - SFROWASA</u>						
N 019618	002 7645801	Jul 24, 2027	DS DP			
<u>MESALAMINE - CANASA</u>						
N 021252	002 8217083	Jun 06, 2028	DP			
	8436051	Jun 06, 2028	DP			
<u>MESALAMINE - APRISO</u>						
N 022301	001 8865688	May 01, 2030		U-1310		
<u>METAXALONE - SKELAXIN</u>						
N 013217	003 7122566	Feb 06, 2026		U-915		
<u>METAXALONE - METAXALONE</u>						
N 022503	001 11918559	Jul 29, 2039	DP			
<u>METFORMIN HYDROCHLORIDE - GLUMETZA</u>						
N 021748	002 7780987	Mar 23, 2025	DS DP			
<u>METFORMIN HYDROCHLORIDE - RIOMET ER</u>						
N 212595	001 9962336	May 01, 2035	DP			
<u>METFORMIN HYDROCHLORIDE; PIOGLITAZONE HYDROCHLORIDE - ACTOPLUS MET</u>						
N 021842	001 9101660	Jan 22, 2027	DP			
	9320714	Feb 03, 2029	DP			
<u>METFORMIN HYDROCHLORIDE; PIOGLITAZONE HYDROCHLORIDE - ACTOPLUS MET</u>						
N 021842	002 9101660	Jan 22, 2027	DP			
	9320714	Feb 03, 2029	DP			
<u>METFORMIN HYDROCHLORIDE; PIOGLITAZONE HYDROCHLORIDE - ACTOPLUS MET XR</u>						
N 022024	001 7785627	Jul 31, 2026	DP			
	7959946	Jul 31, 2026	DP			
<u>METFORMIN HYDROCHLORIDE; PIOGLITAZONE HYDROCHLORIDE - ACTOPLUS MET XR</u>						
N 022024	002 7785627	Jul 31, 2026	DP			
	7959946	Jul 31, 2026	DP			
<u>METFORMIN HYDROCHLORIDE; SAXAGLIPTIN HYDROCHLORIDE - KOMBIGLYZE XR</u>						
N 200678	001 8628799	Jul 13, 2025	DP			
	9339472	Jul 13, 2025	DP			
<u>METFORMIN HYDROCHLORIDE; SAXAGLIPTIN HYDROCHLORIDE - KOMBIGLYZE XR</u>						
N 200678	002 8628799	Jul 13, 2025	DP			
	9339472	Jul 13, 2025	DP			
<u>METFORMIN HYDROCHLORIDE; SAXAGLIPTIN HYDROCHLORIDE - KOMBIGLYZE XR</u>						
N 200678	003 8628799	Jul 13, 2025	DP			
	9339472	Jul 13, 2025	DP			
<u>METFORMIN HYDROCHLORIDE; SITAGLIPTIN PHOSPHATE - JANUMET</u>						
N 022044	001 7326708	Nov 24, 2026	DS DP	U-802		
	7326708*PED	May 24, 2027				
	8414921	Jul 21, 2028	DP	U-1036		
	8414921*PED	Jan 21, 2029				
<u>METFORMIN HYDROCHLORIDE; SITAGLIPTIN PHOSPHATE - JANUMET</u>						
N 022044	002 7326708	Nov 24, 2026	DS DP	U-802		
	7326708*PED	May 24, 2027				
	8414921	Jul 21, 2028	DP	U-1036		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>METFORMIN HYDROCHLORIDE; SITAGLIPTIN PHOSPHATE - JANUMET</u>						
N 022044	002	8414921*PED	Jan 21, 2029			
<u>METFORMIN HYDROCHLORIDE; SITAGLIPTIN PHOSPHATE - JANUMET XR</u>						
N 202270	001	7326708	Nov 24, 2026	DS DP U-1227		
		7326708*PED	May 24, 2027			
<u>METFORMIN HYDROCHLORIDE; SITAGLIPTIN PHOSPHATE - JANUMET XR</u>						
N 202270	002	7326708	Nov 24, 2026	DS DP U-1227		
		7326708*PED	May 24, 2027			
<u>METFORMIN HYDROCHLORIDE; SITAGLIPTIN PHOSPHATE - JANUMET XR</u>						
N 202270	003	7326708	Nov 24, 2026	DS DP U-1227		
		7326708*PED	May 24, 2027			
<u>METHOHEXITAL SODIUM - METHOHEXITAL SODIUM</u>						
A 215488	001				CGT	Jun 10, 2025
<u>METHOTREXATE - OTREXUP</u>						
N 204824	001	10709844	Mar 10, 2029	DP		
		11446441	Jan 24, 2026	DP		
		11497753	Mar 19, 2030	DP		
		11684723	Mar 10, 2029	DP		
		8021335	Oct 04, 2026	DP		
		8480631	Mar 19, 2030	DP U-1442		
		8562564	Jan 24, 2026	DP		
		8579865	Mar 19, 2030	DP U-1442		
		8814834	May 27, 2031	DP		
		8945063	Mar 19, 2030	DP U-1442		
		9421333	Mar 19, 2030	DP U-1442		
		9533102	Jan 24, 2026	DP		
		9629959	Jan 24, 2026	DP		
		9867949	Mar 10, 2029	DP		
<u>METHOTREXATE - OTREXUP</u>						
N 204824	002	10709844	Mar 10, 2029	DP		
		11446441	Jan 24, 2026	DP		
		11497753	Mar 19, 2030	DP		
		11684723	Mar 10, 2029	DP		
		8021335	Oct 04, 2026	DP		
		8480631	Mar 19, 2030	DP U-1442		
		8562564	Jan 24, 2026	DP		
		8579865	Mar 19, 2030	DP U-1442		
		8814834	May 27, 2031	DP		
		8945063	Mar 19, 2030	DP U-1442		
		9421333	Mar 19, 2030	DP U-1442		
		9533102	Jan 24, 2026	DP		
		9629959	Jan 24, 2026	DP		
		9867949	Mar 10, 2029	DP		
<u>METHOTREXATE - OTREXUP</u>						
N 204824	003	10709844	Mar 10, 2029	DP		
		11446441	Jan 24, 2026	DP		
		11497753	Mar 19, 2030	DP		
		11684723	Mar 10, 2029	DP		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>METHOTREXATE - OTREXUP</u>						
N 204824 003	8021335	Oct 04, 2026	DP			
	8480631	Mar 19, 2030	DP U-1442			
	8562564	Jan 24, 2026	DP			
	8579865	Mar 19, 2030	DP U-1442			
	8814834	May 27, 2031	DP			
	8945063	Mar 19, 2030	DP U-1442			
	9421333	Mar 19, 2030	DP U-1442			
	9533102	Jan 24, 2026	DP			
	9629959	Jan 24, 2026	DP			
	9867949	Mar 10, 2029	DP			
<u>METHOTREXATE - OTREXUP</u>						
N 204824 004	10709844	Mar 10, 2029	DP			
	11446441	Jan 24, 2026	DP			
	11497753	Mar 19, 2030	DP			
	11684723	Mar 10, 2029	DP			
	8021335	Oct 04, 2026	DP			
	8480631	Mar 19, 2030	DP U-1442			
	8562564	Jan 24, 2026	DP			
	8579865	Mar 19, 2030	DP U-1442			
	8814834	May 27, 2031	DP			
	8945063	Mar 19, 2030	DP U-1442			
	9421333	Mar 19, 2030	DP U-1442			
	9533102	Jan 24, 2026	DP			
	9629959	Jan 24, 2026	DP			
	9867949	Mar 10, 2029	DP			
<u>METHOTREXATE - OTREXUP</u>						
N 204824 005	10709844	Mar 10, 2029	DP			
	11446441	Jan 24, 2026	DP			
	11497753	Mar 19, 2030	DP			
	11684723	Mar 10, 2029	DP			
	8021335	Oct 04, 2026	DP			
	8480631	Mar 19, 2030	DP U-1442			
	8562564	Jan 24, 2026	DP			
	8579865	Mar 19, 2030	DP U-1442			
	8814834	May 27, 2031	DP			
	8945063	Mar 19, 2030	DP U-1442			
	9421333	Mar 19, 2030	DP U-1442			
	9533102	Jan 24, 2026	DP			
	9629959	Jan 24, 2026	DP			
	9867949	Mar 10, 2029	DP			
<u>METHOTREXATE - OTREXUP</u>						
N 204824 006	10709844	Mar 10, 2029	DP			
	11446441	Jan 24, 2026	DP			
	11497753	Mar 19, 2030	DP			
	11684723	Mar 10, 2029	DP			
	8021335	Oct 04, 2026	DP			
	8480631	Mar 19, 2030	DP U-1442			
	8562564	Jan 24, 2026	DP			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>METHOTREXATE - OTREXUP</u>						
N 204824	006	8579865	Mar 19, 2030	DP U-1442		
		8814834	May 27, 2031	DP		
		8945063	Mar 19, 2030	DP U-1442		
		9421333	Mar 19, 2030	DP U-1442		
		9533102	Jan 24, 2026	DP		
		9629959	Jan 24, 2026	DP		
		9867949	Mar 10, 2029	DP		
<u>METHOTREXATE - OTREXUP</u>						
N 204824	007	10709844	Mar 10, 2029	DP		
		11446441	Jan 24, 2026	DP		
		11497753	Mar 19, 2030	DP		
		11684723	Mar 10, 2029	DP		
		8021335	Oct 04, 2026	DP		
		8480631	Mar 19, 2030	DP U-1442		
		8562564	Jan 24, 2026	DP		
		8579865	Mar 19, 2030	DP U-1442		
		8814834	May 27, 2031	DP		
		8945063	Mar 19, 2030	DP U-1442		
		9421333	Mar 19, 2030	DP U-1442		
		9533102	Jan 24, 2026	DP		
		9629959	Jan 24, 2026	DP		
		9867949	Mar 10, 2029	DP		
<u>METHOTREXATE - OTREXUP</u>						
N 204824	008	10709844	Mar 10, 2029	DP		
		11446441	Jan 24, 2026	DP		
		11497753	Mar 19, 2030	DP		
		11684723	Mar 10, 2029	DP		
		8021335	Oct 04, 2026	DP		
		8480631	Mar 19, 2030	DP U-1442		
		8562564	Jan 24, 2026	DP		
		8579865	Mar 19, 2030	DP U-1442		
		8814834	May 27, 2031	DP		
		8945063	Mar 19, 2030	DP U-1442		
		9421333	Mar 19, 2030	DP U-1442		
		9533102	Jan 24, 2026	DP		
		9629959	Jan 24, 2026	DP		
		9867949	Mar 10, 2029	DP		
<u>METHOTREXATE - RASUVO</u>						
N 205776	001	8664231	Jun 01, 2029	U-1442		
<u>METHOTREXATE - RASUVO</u>						
N 205776	002	8664231	Jun 01, 2029	U-1442		
<u>METHOTREXATE - RASUVO</u>						
N 205776	003	8664231	Jun 01, 2029	U-1442		
<u>METHOTREXATE - RASUVO</u>						
N 205776	004	8664231	Jun 01, 2029	U-1442		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>METHOTREXATE - RASUVO</u>						
N 205776	005 8664231	Jun 01, 2029	U-1442			
<u>METHOTREXATE - RASUVO</u>						
N 205776	006 8664231	Jun 01, 2029	U-1442			
<u>METHOTREXATE - RASUVO</u>						
N 205776	007 8664231	Jun 01, 2029	U-1442			
<u>METHOTREXATE - RASUVO</u>						
N 205776	008 8664231	Jun 01, 2029	U-1442			
<u>METHOTREXATE - RASUVO</u>						
N 205776	009 8664231	Jun 01, 2029	U-1442			
<u>METHOTREXATE - RASUVO</u>						
N 205776	010 8664231	Jun 01, 2029	U-1442			
<u>METHOTREXATE - JYLAMVO</u>						
N 212479	001 11129833	Oct 28, 2035	DP			
	11771701	Oct 29, 2034	DP U-3700			
	11771701	Oct 29, 2034	DP U-3701			
	11771701	Oct 29, 2034	DP U-3702			
	11771701	Oct 29, 2034	DP U-3703			
	11771701	Oct 29, 2034	DP U-3704			
<u>METHOTREXATE SODIUM - XATMEP</u>						
N 208400	001 10231927	Jan 02, 2033	U-1349			
	10231927	Jan 02, 2033	U-1699			
	10610485	Jan 02, 2033	DP			
	11116724	Jan 02, 2033	U-1349			
	11116724	Jan 02, 2033	U-1699			
	11969503	Jan 02, 2033	DP			
	9259427	Jan 02, 2033	DP			
	9855215	Jan 02, 2033	DP			
<u>METHYLENE BLUE - PROVAYBLUE</u>						
N 204630	001				M-303	Jan 08, 2027
<u>METHYLENE BLUE - PROVAYBLUE</u>						
N 204630	002				M-303	Jan 08, 2027
<u>METHYLNALTREXONE BROMIDE - RELISTOR</u>						
N 021964	001 8247425	Dec 31, 2030	U-1185			
	8420663	Sep 30, 2029	U-1185			
	8822490	Sep 30, 2029	DP U-1185			
	9180125	Sep 30, 2029	DP U-1185			
	9492445	Sep 30, 2029	DP U-1185			
<u>METHYLNALTREXONE BROMIDE - RELISTOR</u>						
N 021964	002 8247425	Dec 31, 2030	U-1185			
	8420663	Sep 30, 2029	U-1185			
	8822490	Sep 30, 2029	DP U-1185			
	9180125	Sep 30, 2029	DP U-1185			
	9492445	Sep 30, 2029	DP U-1185			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>METHYLNALTREXONE BROMIDE - RELISTOR</u>						
N 021964	003	8247425	Dec 31, 2030			U-1185
		8420663	Sep 30, 2029			U-1185
		8822490	Sep 30, 2029	DP		U-1185
		9180125	Sep 30, 2029	DP		U-1185
		9492445	Sep 30, 2029	DP		U-1185
<u>METHYLNALTREXONE BROMIDE - RELISTOR</u>						
N 208271	001	10307417	Mar 10, 2031			DP
		10376505	Mar 10, 2031			DP
		8420663	Sep 30, 2029			U-1185
		8524276	Mar 10, 2031			DP
		8956651	Mar 10, 2031			DP
		9180125	Sep 30, 2029	DP		U-1185
		9314461	Mar 10, 2031			DP
		9492445	Sep 30, 2029	DP		U-1185
		9724343	Sep 30, 2029	DP		U-1185
<u>METHYLPHENIDATE - DAYTRANA</u>						
N 021514	001	8632802	Oct 07, 2025			DP
		9034370	Oct 07, 2025			DP
		9668981	Oct 07, 2025			U-2024
<u>METHYLPHENIDATE - DAYTRANA</u>						
N 021514	002	8632802	Oct 07, 2025			DP
		9034370	Oct 07, 2025			DP
		9668981	Oct 07, 2025			U-2024
<u>METHYLPHENIDATE - DAYTRANA</u>						
N 021514	003	8632802	Oct 07, 2025			DP
		9034370	Oct 07, 2025			DP
		9668981	Oct 07, 2025			U-2024
<u>METHYLPHENIDATE - DAYTRANA</u>						
N 021514	004	8632802	Oct 07, 2025			DP
		9034370	Oct 07, 2025			DP
		9668981	Oct 07, 2025			U-2024
<u>METHYLPHENIDATE - COTEMPLA XR-ODT</u>						
N 205489	001	11166947	Jan 25, 2038			U-3299
		8840924	Jun 05, 2026			DP
		9072680	Jun 28, 2032			DP
		9089496	Jun 28, 2032			DP
<u>METHYLPHENIDATE - COTEMPLA XR-ODT</u>						
N 205489	002	11166947	Jan 25, 2038			U-3299
		8840924	Jun 05, 2026			DP
		9072680	Jun 28, 2032			DP
		9089496	Jun 28, 2032			DP
<u>METHYLPHENIDATE - COTEMPLA XR-ODT</u>						
N 205489	003	11166947	Jan 25, 2038			U-3299
		8840924	Jun 05, 2026			DP
		9072680	Jun 28, 2032			DP
		9089496	Jun 28, 2032			DP

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>METHYLPHENIDATE HYDROCHLORIDE - QUILLIVANT XR</u>						
N 202100 001	8062667	Mar 29, 2029	DP			
	8287903	Feb 15, 2031	DP			
	8465765	Feb 15, 2031	DP U-1415			
	8563033	Feb 15, 2031	DP U-1415			
	8778390	Feb 15, 2031	DP U-1543			
	8956649	Feb 15, 2031	DP U-1665			
	9040083	Feb 15, 2031	DP			
<u>METHYLPHENIDATE HYDROCHLORIDE - QUILLICHEW ER</u>						
N 207960 001	10857143	Aug 14, 2033	DP U-2993			
	11103494	Aug 14, 2033	DP			
	11103495	Aug 14, 2033	DP U-2993			
	11633389	Aug 14, 2033	DP			
	8202537	Mar 15, 2027	DP			
	8287903	Feb 15, 2031	DP			
	8999386	Aug 14, 2033	DP			
	9295642	Aug 14, 2033	DP U-1827			
	9545399	Aug 14, 2033	DP U-1827			
	9844544	Aug 14, 2033	DP U-2203			
<u>METHYLPHENIDATE HYDROCHLORIDE - QUILLICHEW ER</u>						
N 207960 002	10857143	Aug 14, 2033	DP U-2993			
	11103494	Aug 14, 2033	DP			
	11103495	Aug 14, 2033	DP U-2993			
	11633389	Aug 14, 2033	DP			
	8202537	Mar 15, 2027	DP			
	8287903	Feb 15, 2031	DP			
	8999386	Aug 14, 2033	DP			
	9295642	Aug 14, 2033	DP U-1827			
	9545399	Aug 14, 2033	DP U-1827			
	9844544	Aug 14, 2033	DP U-2203			
<u>METHYLPHENIDATE HYDROCHLORIDE - QUILLICHEW ER</u>						
N 207960 003	10857143	Aug 14, 2033	DP U-2993			
	11103494	Aug 14, 2033	DP			
	11103495	Aug 14, 2033	DP U-2993			
	11633389	Aug 14, 2033	DP			
	8202537	Mar 15, 2027	DP			
	8287903	Feb 15, 2031	DP			
	8999386	Aug 14, 2033	DP			
	9295642	Aug 14, 2033	DP U-1827			
	9545399	Aug 14, 2033	DP U-1827			
	9844544	Aug 14, 2033	DP U-2203			
<u>METHYLPHENIDATE HYDROCHLORIDE - JORNAY PM</u>						
N 209311 001	10182995	Mar 23, 2032	DP			
	10292937	Mar 23, 2032	U-2357			
	10617651	Mar 23, 2032	U-2357			
	10881618	Mar 23, 2032	U-2357			
	10905652	Mar 23, 2032	DP			
	11241391	Mar 23, 2032	U-2357			
	11241392	Mar 23, 2032	DP			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>METHYLPHENIDATE HYDROCHLORIDE - JORNAY PM</u>						
N 209311 001	11911518	Mar 23, 2032				
	8916588	Mar 23, 2032				
	8927010	Mar 23, 2032	DP			
	9023389	Mar 23, 2032	DP			
	9028868	Mar 23, 2032				
	9034902	Mar 23, 2032				
	9283214	Mar 23, 2032	DP			
	9498447	Mar 23, 2032	DP			
	9603809	Mar 23, 2032				
<u>METHYLPHENIDATE HYDROCHLORIDE - JORNAY PM</u>						
N 209311 002	10182995	Mar 23, 2032				
	10292937	Mar 23, 2032				
	10617651	Mar 23, 2032				
	10881618	Mar 23, 2032				
	10905652	Mar 23, 2032	DP			
	11241391	Mar 23, 2032				
	11241392	Mar 23, 2032	DP			
	11911518	Mar 23, 2032				
	8916588	Mar 23, 2032				
	8927010	Mar 23, 2032	DP			
	9023389	Mar 23, 2032	DP			
	9028868	Mar 23, 2032				
	9034902	Mar 23, 2032				
	9283214	Mar 23, 2032	DP			
	9498447	Mar 23, 2032	DP			
	9603809	Mar 23, 2032				
<u>METHYLPHENIDATE HYDROCHLORIDE - JORNAY PM</u>						
N 209311 003	10182995	Mar 23, 2032				
	10292937	Mar 23, 2032				
	10617651	Mar 23, 2032				
	10881618	Mar 23, 2032				
	10905652	Mar 23, 2032	DP			
	11241391	Mar 23, 2032				
	11241392	Mar 23, 2032	DP			
	11911518	Mar 23, 2032				
	8916588	Mar 23, 2032				
	8927010	Mar 23, 2032	DP			
	9023389	Mar 23, 2032	DP			
	9028868	Mar 23, 2032				
	9034902	Mar 23, 2032				
	9283214	Mar 23, 2032	DP			
	9498447	Mar 23, 2032	DP			
	9603809	Mar 23, 2032				
<u>METHYLPHENIDATE HYDROCHLORIDE - JORNAY PM</u>						
N 209311 004	10182995	Mar 23, 2032				
	10292937	Mar 23, 2032				
	10617651	Mar 23, 2032				
	10881618	Mar 23, 2032				

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>METHYLPHENIDATE HYDROCHLORIDE - JORNAY PM</u>						
N 209311 004	10905652	Mar 23, 2032	DP			
	11241391	Mar 23, 2032		U-2357		
	11241392	Mar 23, 2032	DP			
	11911518	Mar 23, 2032		U-2357		
	8916588	Mar 23, 2032		U-2357		
	8927010	Mar 23, 2032	DP			
	9023389	Mar 23, 2032	DP			
	9028868	Mar 23, 2032		U-2357		
	9034902	Mar 23, 2032		U-2357		
	9283214	Mar 23, 2032	DP			
	9498447	Mar 23, 2032	DP			
	9603809	Mar 23, 2032		U-2357		
<u>METHYLPHENIDATE HYDROCHLORIDE - JORNAY PM</u>						
N 209311 005	10182995	Mar 23, 2032	DP			
	10292937	Mar 23, 2032		U-2357		
	10617651	Mar 23, 2032		U-2357		
	10881618	Mar 23, 2032		U-2357		
	10905652	Mar 23, 2032	DP			
	11241391	Mar 23, 2032		U-2357		
	11241392	Mar 23, 2032	DP			
	11911518	Mar 23, 2032		U-2357		
	8916588	Mar 23, 2032		U-2357		
	8927010	Mar 23, 2032	DP			
	9023389	Mar 23, 2032	DP			
	9028868	Mar 23, 2032		U-2357		
	9034902	Mar 23, 2032		U-2357		
	9283214	Mar 23, 2032	DP			
	9498447	Mar 23, 2032	DP			
	9603809	Mar 23, 2032		U-2357		
<u>METHYLPHENIDATE HYDROCHLORIDE - ADHANSIA XR</u>						
N 212038 001	10111839	Oct 30, 2035		U-2357		
	10292938	Oct 30, 2035	DP			
	10292939	Oct 30, 2035	DP	U-2357		
	10449159	Oct 30, 2035		U-2357		
	10500162	Oct 30, 2035		U-2357		
	10507186	Oct 30, 2035	DP			
	10512612	Oct 30, 2035	DP			
	10512613	Oct 30, 2035		U-2357		
	10568841	Oct 30, 2035	DP	U-2357		
	10688060	Oct 30, 2035	DP			
	10722473	Nov 19, 2038		U-2357		
	9974752	Oct 30, 2035	DP			
<u>METHYLPHENIDATE HYDROCHLORIDE - ADHANSIA XR</u>						
N 212038 002	10111839	Oct 30, 2035		U-2357		
	10292938	Oct 30, 2035	DP			
	10292939	Oct 30, 2035	DP	U-2357		
	10449159	Oct 30, 2035		U-2357		
	10500162	Oct 30, 2035		U-2357		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>METHYLPHENIDATE HYDROCHLORIDE - ADHANSIA XR</u>						
N 212038 002	10507186	Oct 30, 2035	DP			
	10512612	Oct 30, 2035	DP			
	10512613	Oct 30, 2035		U-2357		
	10568841	Oct 30, 2035	DP	U-2357		
	10688060	Oct 30, 2035	DP			
	10722473	Nov 19, 2038		U-2357		
	9974752	Oct 30, 2035	DP			
<u>METHYLPHENIDATE HYDROCHLORIDE - ADHANSIA XR</u>						
N 212038 003	10111839	Oct 30, 2035		U-2357		
	10292938	Oct 30, 2035	DP			
	10292939	Oct 30, 2035	DP	U-2357		
	10449159	Oct 30, 2035		U-2357		
	10500162	Oct 30, 2035		U-2357		
	10507186	Oct 30, 2035	DP			
	10512612	Oct 30, 2035	DP			
	10512613	Oct 30, 2035		U-2357		
	10568841	Oct 30, 2035	DP	U-2357		
	10688060	Oct 30, 2035	DP			
	10722473	Nov 19, 2038		U-2357		
	9974752	Oct 30, 2035	DP			
<u>METHYLPHENIDATE HYDROCHLORIDE - ADHANSIA XR</u>						
N 212038 004	10111839	Oct 30, 2035		U-2357		
	10292938	Oct 30, 2035	DP			
	10292939	Oct 30, 2035	DP	U-2357		
	10449159	Oct 30, 2035		U-2357		
	10500162	Oct 30, 2035		U-2357		
	10507186	Oct 30, 2035	DP			
	10512612	Oct 30, 2035	DP			
	10512613	Oct 30, 2035		U-2357		
	10568841	Oct 30, 2035	DP	U-2357		
	10688060	Oct 30, 2035	DP			
	10722473	Nov 19, 2038		U-2357		
	9974752	Oct 30, 2035	DP			
<u>METHYLPHENIDATE HYDROCHLORIDE - ADHANSIA XR</u>						
N 212038 005	10111839	Oct 30, 2035		U-2357		
	10292938	Oct 30, 2035	DP			
	10292939	Oct 30, 2035	DP	U-2357		
	10449159	Oct 30, 2035		U-2357		
	10500162	Oct 30, 2035		U-2357		
	10507186	Oct 30, 2035	DP			
	10512612	Oct 30, 2035	DP			
	10512613	Oct 30, 2035		U-2357		
	10568841	Oct 30, 2035	DP	U-2357		
	10688060	Oct 30, 2035	DP			
	10722473	Nov 19, 2038		U-2357		
	9974752	Oct 30, 2035	DP			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>METHYLPHENIDATE HYDROCHLORIDE - ADHANSIA XR</u>						
N 212038 006	10111839	Oct 30, 2035	U-2357			
	10292938	Oct 30, 2035	DP			
	10292939	Oct 30, 2035	DP U-2357			
	10449159	Oct 30, 2035	U-2357			
	10500162	Oct 30, 2035	U-2357			
	10507186	Oct 30, 2035	DP			
	10512612	Oct 30, 2035	DP			
	10512613	Oct 30, 2035	U-2357			
	10568841	Oct 30, 2035	DP U-2357			
	10688060	Oct 30, 2035	DP			
	10722473	Nov 19, 2038	U-2357			
	9974752	Oct 30, 2035	DP			
<u>METHYLPHENIDATE HYDROCHLORIDE - RELEXXII</u>						
N 216117 001	10265308	Feb 03, 2037	DP U-4032			
	10695336	Feb 03, 2037	U-4032			
	9707217	Feb 03, 2037	DP			
	9827234	Feb 03, 2037	U-4032			
	9855258	Feb 03, 2037	DP U-4032			
<u>METHYLPHENIDATE HYDROCHLORIDE - RELEXXII</u>						
N 216117 002	10265308	Feb 03, 2037	DP U-4032			
	10695336	Feb 03, 2037	U-4032			
	9707217	Feb 03, 2037	DP			
	9827234	Feb 03, 2037	U-4032			
	9855258	Feb 03, 2037	DP U-4032			
<u>METHYLPHENIDATE HYDROCHLORIDE - RELEXXII</u>						
N 216117 003	10265308	Feb 03, 2037	DP U-4032			
	10695336	Feb 03, 2037	U-4032			
	9707217	Feb 03, 2037	DP			
	9827234	Feb 03, 2037	U-4032			
	9855258	Feb 03, 2037	DP U-4032			
<u>METHYLPHENIDATE HYDROCHLORIDE - RELEXXII</u>						
N 216117 004	10265308	Feb 03, 2037	DP U-4032			
	10695336	Feb 03, 2037	U-4032			
	9707217	Feb 03, 2037	DP			
	9827234	Feb 03, 2037	U-4032			
	9855258	Feb 03, 2037	DP U-4032			
<u>METHYLPHENIDATE HYDROCHLORIDE - RELEXXII</u>						
N 216117 005	10265308	Feb 03, 2037	DP U-4032			
	10695336	Feb 03, 2037	U-4032			
	9707217	Feb 03, 2037	DP			
	9827234	Feb 03, 2037	U-4032			
	9855258	Feb 03, 2037	DP U-4032			
<u>METHYLPHENIDATE HYDROCHLORIDE - RELEXXII</u>						
N 216117 006	10265308	Feb 03, 2037	DP U-4032			
	10695336	Feb 03, 2037	U-4032			
	9707217	Feb 03, 2037	DP			
	9827234	Feb 03, 2037	U-4032			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>METHYLPHENIDATE HYDROCHLORIDE - RELEXXII</u>						
N 216117	006	9855258	Feb 03, 2037	DP U-4032		
<u>METHYLPHENIDATE HYDROCHLORIDE - RELEXXII</u>						
N 216117	007	10265308	Feb 03, 2037	DP U-4032		
		10695336	Feb 03, 2037	U-4032		
		9707217	Feb 03, 2037	DP		
		9827234	Feb 03, 2037	U-4032		
		9855258	Feb 03, 2037	DP U-4032		
<u>METOCLOPRAMIDE HYDROCHLORIDE - GIMOTI</u>						
N 209388	001	11020361	Dec 22, 2029	U-2843		
		11628150	Dec 22, 2029	DP U-2843		
		11813231	Dec 22, 2029	DP U-2843		
		8334281	May 16, 2030	DP U-2843		
<u>METOPROLOL SUCCINATE - KAPSPARGO SPRINKLE</u>						
N 210428	001	9504655	Jul 09, 2035	DP		
		9700530	Jul 09, 2035	DP		
<u>METOPROLOL SUCCINATE - KAPSPARGO SPRINKLE</u>						
N 210428	002	9504655	Jul 09, 2035	DP		
		9700530	Jul 09, 2035	DP		
<u>METOPROLOL SUCCINATE - KAPSPARGO SPRINKLE</u>						
N 210428	003	9504655	Jul 09, 2035	DP		
		9700530	Jul 09, 2035	DP		
<u>METOPROLOL SUCCINATE - KAPSPARGO SPRINKLE</u>						
N 210428	004	9504655	Jul 09, 2035	DP		
		9700530	Jul 09, 2035	DP		
<u>METRONIDAZOLE - NUVESSA</u>						
N 205223	001	10238634	Jun 28, 2032	DP		
		10596155	Jun 28, 2032	DP		
		7893097	Feb 19, 2028	DP		
		8658678	Jun 27, 2028	U-1682		
		8877792	Feb 02, 2028	DP		
		8946276	Jun 28, 2032	U-1664		
		9198858	Jun 28, 2032	U-1664		
<u>METRONIDAZOLE - LIKMEZ</u>						
N 216755	001	11541035	Oct 04, 2039	DP		
<u>MICONAZOLE NITRATE; WHITE PETROLATUM; ZINC OXIDE - VUSION</u>						
N 021026	001	8147852	Mar 30, 2028	U-1426		
<u>MIDAZOLAM - NAYZILAM</u>						
N 211321	001	8217033	Jan 18, 2028	DP U-2526	ODE-243	May 17, 2026
		8809322	Jan 18, 2028	DP		
		9289432	Jan 18, 2028	DP U-2526		
		9687495	Jan 18, 2028	DP U-2526		
<u>MIDAZOLAM - MIDAZOLAM IN 0.9% SODIUM CHLORIDE</u>						
N 211844	001	10966990	Jun 20, 2038	DP		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>MIDAZOLAM - MIDAZOLAM IN 0.9% SODIUM CHLORIDE</u>						
N 211844	002 10966990	Jun 20, 2038	DP			
<u>MIDAZOLAM HYDROCHLORIDE - SEIZALAM</u>						
N 209566	001				ODE-207	Sep 14, 2025
<u>MIDOSTAURIN - RYDAPT</u>						
N 207997	001 7973031	Oct 09, 2028	U-2007			
	8575146	Dec 02, 2030	U-2008			
<u>MIFEPRISTONE - KORLYM</u>						
N 202107	001 10006924	Aug 12, 2036	U-1643			
	10151763	Jan 18, 2037	U-1643			
	10166242	Apr 20, 2036	U-1643			
	10166243	Apr 20, 2036	U-1643			
	10195214	Jun 19, 2037	U-1643			
	10231983	Aug 22, 2038	U-1643			
	10314850	Aug 22, 2038	U-1643			
	10495650	Aug 12, 2036	U-1643			
	10500216	Mar 05, 2033	U-1643			
	10660904	Apr 20, 2036	U-1643			
	10780097	Aug 22, 2038	U-1643			
	10842800	Jun 19, 2037	U-1643			
	10842801	Nov 15, 2032	U-1643			
	11969435	Jun 19, 2037	U-1643			
	12097210	Jun 19, 2037	U-1643			
	8921348	Aug 27, 2028	U-1643			
	9829495	Aug 15, 2036	U-1643			
	9943526	Apr 20, 2036	U-1643			
<u>MIGALASTAT HYDROCHLORIDE - GALAFOLD</u>						
N 208623	001 10076514	Mar 15, 2037	U-2371		ODE-205	Aug 10, 2025
	10251873	May 30, 2038	U-2371			
	10383864	May 16, 2027	U-2371			
	10406143	May 16, 2027	U-2371			
	10471053	May 30, 2038	U-2371			
	10525045	Apr 28, 2028	U-2371			
	10792278	May 30, 2038	U-2371			
	10792279	May 30, 2038	U-2371			
	10799491	May 30, 2038	U-2371			
	10806727	May 30, 2038	U-2371			
	10813921	Feb 12, 2029	U-2371			
	10849889	May 30, 2038	U-2371			
	10849890	May 30, 2038	U-2371			
	10857141	May 30, 2038	U-2371			
	10857142	May 30, 2038	U-2371			
	10874655	May 30, 2038	U-2371			
	10874656	May 30, 2038	U-2371			
	10874657	May 30, 2038	U-2371			
	10925866	Apr 28, 2028	U-2371			
	11033538	Apr 28, 2028	U-2371			
	11234972	Mar 15, 2037	U-2371			
	11241422	May 16, 2027	U-2371			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>MIGALASTAT HYDROCHLORIDE - GALAFOLD</u>						
N 208623	001	11278536	May 30, 2038	U-2371		
		11278537	May 30, 2038	U-2371		
		11278538	May 30, 2038	U-2371		
		11278539	May 30, 2038	U-2371		
		11278540	May 30, 2038	U-2371		
		11304940	May 30, 2038	DS		
		11357761	May 30, 2038	U-2371		
		11357762	May 30, 2038	U-2371		
		11357763	May 30, 2038	U-2371		
		11357764	May 30, 2038	DS		
		11357765	May 30, 2038	DS		
		11357784	Feb 06, 2039	U-2371		
		11376244	May 30, 2038	DS		
		11389436	May 30, 2038	U-2371		
		11389437	May 30, 2038	U-2371		
		11426396	May 30, 2038	DS		
		11458128	May 30, 2038	U-2371		
		11612593	May 30, 2038	DS		
		11612594	May 30, 2038	DS		
		11622962	Mar 17, 2039	DS		
		11633387	May 30, 2038	DS		
		11633388	Mar 25, 2039	U-2371		
		11642334	Feb 20, 2039	U-2371		
		11666564	May 30, 2038	U-2371		
		11786516	May 30, 2038	DS		
		11813255	May 30, 2038	U-2371		
		11826360	Feb 16, 2039	DS		
		11833164	Jan 11, 2042	U-2371		
		11903938	Aug 17, 2038	DS U-2371		
		12042488	May 30, 2038	DS		
		12042489	May 30, 2038	DS U-2371		
		12042490	May 30, 2038	U-2371		
		12109205	May 30, 2038	DS U-2371		
		8592362	Feb 12, 2029	U-2371	Y	
		9000011	May 16, 2027	U-2371		
		9095584	Feb 12, 2029	U-2371		
		9480682	May 16, 2027	U-2371		
		9987263	May 16, 2027	U-2371		
		9999618	Apr 28, 2028	U-2372		
		9999618	Apr 28, 2028	U-2373		
		RE48608	Oct 20, 2031	U-2371		
<u>MIGLUSTAT - OPFOLDA</u>						
N 215211	001	10208299	Sep 30, 2035	DS DP U-3726	NP	Sep 28, 2026
		10512677	Mar 07, 2033	U-3726		
		10857212	Aug 12, 2037	U-3726		
		10961522	Sep 30, 2035	U-3726		
		11278599	Mar 07, 2033	U-3726		
		11278601	Dec 29, 2036	DP U-3726		
		11753632	Sep 30, 2035	U-3726		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>MIGLUSTAT - OPFOLDA</u>						
N 215211	001	10208299	Sep 30, 2035	DS DP U-3726	NP	Sep 28, 2026
		10512677	Mar 07, 2033	U-3726		
		10857212	Aug 12, 2037	U-3726		
		10961522	Sep 30, 2035	U-3726		
		11278599	Mar 07, 2033	U-3726		
		11278601	Dec 29, 2036	DP U-3726		
		11753632	Sep 30, 2035	U-3726		
<u>MILNACIPRAN HYDROCHLORIDE - SAVELLA</u>						
N 022256	001	7994220	Sep 19, 2029	U-819		
<u>MILNACIPRAN HYDROCHLORIDE - SAVELLA</u>						
N 022256	002	7994220	Sep 19, 2029	U-819		
<u>MILNACIPRAN HYDROCHLORIDE - SAVELLA</u>						
N 022256	003	7994220	Sep 19, 2029	U-819		
<u>MILNACIPRAN HYDROCHLORIDE - SAVELLA</u>						
N 022256	004	7994220	Sep 19, 2029	U-819		
<u>MINOCYCLINE HYDROCHLORIDE - MINOCIN</u>						
N 050444	001	11944634	Oct 16, 2032	DP		
		12161656	May 12, 2031	DP		
		9084802	May 12, 2031	U-282		
		9278105	May 12, 2031	U-282		
<u>MINOCYCLINE HYDROCHLORIDE - SOLODYN</u>						
N 050808	001	7790705	Jun 24, 2025	U-1078		
		7919483	Mar 07, 2027	U-1078		
		8252776	Jun 24, 2025	U-124		
		8268804	Jun 24, 2025	U-1078		
<u>MINOCYCLINE HYDROCHLORIDE - SOLODYN</u>						
N 050808	002	7541347	Apr 02, 2027	U-917		
		7544373	Apr 02, 2027	DP		
		7790705	Jun 24, 2025	U-1078		
		7919483	Mar 07, 2027	U-1078		
		8252776	Jun 24, 2025	U-124		
		8268804	Jun 24, 2025	U-1078		
<u>MINOCYCLINE HYDROCHLORIDE - SOLODYN</u>						
N 050808	003	7790705	Jun 24, 2025	U-1078		
		7919483	Mar 07, 2027	U-1078		
		8252776	Jun 24, 2025	U-124		
		8268804	Jun 24, 2025	U-1078		
<u>MINOCYCLINE HYDROCHLORIDE - SOLODYN</u>						
N 050808	004	7790705	Jun 24, 2025	U-1078		
		7919483	Mar 07, 2027	U-1078		
		8252776	Jun 24, 2025	U-124		
		8268804	Jun 24, 2025	U-1078		
		9192615	Nov 17, 2031	DP		
<u>MINOCYCLINE HYDROCHLORIDE - SOLODYN</u>						
N 050808	005	7790705	Jun 24, 2025	U-1078		
		7919483	Mar 07, 2027	U-1078		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>MINOCYCLINE HYDROCHLORIDE - SOLODYN</u>						
N 050808	005	8252776	Jun 24, 2025	U-124		
		8268804	Jun 24, 2025	U-1078		
		9192615	Nov 17, 2031	DP		
<u>MINOCYCLINE HYDROCHLORIDE - SOLODYN</u>						
N 050808	006	7790705	Jun 24, 2025	U-1078		
		7919483	Mar 07, 2027	U-1078		
		8252776	Jun 24, 2025	U-124		
		8268804	Jun 24, 2025	U-1078		
		8722650	Jun 24, 2025	U-1078		
<u>MINOCYCLINE HYDROCHLORIDE - SOLODYN</u>						
N 050808	007	7790705	Jun 24, 2025	U-1078		
		7919483	Mar 07, 2027	U-1078		
		8252776	Jun 24, 2025	U-124		
		8268804	Jun 24, 2025	U-1078		
		8722650	Jun 24, 2025	U-1078		
<u>MINOCYCLINE HYDROCHLORIDE - SOLODYN</u>						
N 050808	008	7790705	Jun 24, 2025	U-1078		
		7919483	Mar 07, 2027	U-1078		
		8252776	Jun 24, 2025	U-124		
		8268804	Jun 24, 2025	U-1078		
		8722650	Jun 24, 2025	U-1078		
<u>MINOCYCLINE HYDROCHLORIDE - XIMINO</u>						
N 201922	001	7541347	Apr 02, 2027	U-917		
		7544373	Apr 02, 2027	DP		
		7790705	Jun 24, 2025	U-124		
		7919483	Mar 07, 2027	U-124		
		8252776	Jun 24, 2025	U-124		
		8268804	Jun 24, 2025	U-124		
<u>MINOCYCLINE HYDROCHLORIDE - XIMINO</u>						
N 201922	003	7541347	Apr 02, 2027	U-917		
		7544373	Apr 02, 2027	DP		
		7790705	Jun 24, 2025	U-124		
		7919483	Mar 07, 2027	U-124		
		8252776	Jun 24, 2025	U-124		
		8268804	Jun 24, 2025	U-124		
<u>MINOCYCLINE HYDROCHLORIDE - XIMINO</u>						
N 201922	005	7541347	Apr 02, 2027	U-917		
		7544373	Apr 02, 2027	DP		
		7790705	Jun 24, 2025	U-124		
		7919483	Mar 07, 2027	U-124		
		8252776	Jun 24, 2025	U-124		
		8268804	Jun 24, 2025	U-124		
<u>MINOCYCLINE HYDROCHLORIDE - MINOLIRA</u>						
N 209269	001	11103517	Apr 07, 2036	DP		
<u>MINOCYCLINE HYDROCHLORIDE - MINOLIRA</u>						
N 209269	002	11103517	Apr 07, 2036	DP		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>MINOCYCLINE HYDROCHLORIDE - AMZEEQ</u>						
N 212379	001 10086080	Oct 01, 2030	U-2647			
	10137200	Oct 01, 2030	U-2647			
	10213512	Oct 01, 2030	DP U-2647			
	10265404	Oct 01, 2030	DP			
	10398641	Sep 08, 2037	U-2647			
	10517882	Oct 01, 2030	U-2647			
	10821187	Oct 01, 2030	U-2647			
	10849847	Sep 08, 2037	U-2647			
	8865139	Oct 01, 2030	DP U-2647			
	8945516	Oct 01, 2030	DP			
	8992896	Oct 01, 2030	DP U-2647			
	9675700	Oct 01, 2030	DP U-2647			
<u>MINOCYCLINE HYDROCHLORIDE - ZILXI</u>						
N 213690	001 10213512	Oct 01, 2030	DP U-1631			
	10265404	Oct 01, 2030	DP			
	10322186	Oct 01, 2030	U-1631			
	10946101	Oct 01, 2030	U-1631			
	12138311	Oct 01, 2030	U-1631			
	8865139	Oct 01, 2030	DP U-1631			
	8945516	Oct 01, 2030	DP			
	8992896	Oct 01, 2030	DP U-1631			
	9675700	Oct 01, 2030	DP U-1631			
<u>MINOCYCLINE HYDROCHLORIDE - EMROSI</u>						
N 219015	001 10905664	Jan 07, 2039	U-4043		NP	Nov 01, 2027
	11191740	Jan 07, 2039	U-4043			
	11364212	Jan 07, 2039	U-4043			
<u>MIPOMERSEN SODIUM - KYNAMRO</u>						
N 203568	001 7511131	Jan 29, 2027	DS			
<u>MIRABEGRON - MYRBETRIO</u>						
N 202611	001 10842780	Sep 28, 2029	DP U-2996			
	10842780	Sep 28, 2029	DP U-3670			
	10842780*PED	Mar 28, 2030				
	11707451	Sep 28, 2029	U-2996			
	11707451	Sep 28, 2029	U-3670			
	11707451*PED	Mar 28, 2030				
	12059409	Sep 28, 2029	DP			
	12097189	Sep 28, 2029	U-2996			
	12097189	Sep 28, 2029	U-3670			
	8772315	Oct 30, 2028	U-2300			
	8772315*PED	Apr 30, 2029				
<u>MIRABEGRON - MYRBETRIO</u>						
N 202611	002 10842780	Sep 28, 2029	DP U-2996			
	10842780	Sep 28, 2029	DP U-3670			
	10842780*PED	Mar 28, 2030				
	11707451	Sep 28, 2029	U-2996			
	11707451	Sep 28, 2029	U-3670			
	11707451*PED	Mar 28, 2030				
	12059409	Sep 28, 2029	DP			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>MIRABEGRON - MYRBETRIQ</u>						
N 202611 002	12097189	Sep 28, 2029	U-2996			
	12097189	Sep 28, 2029	U-3670			
	8772315	Oct 30, 2028	U-2300			
	8772315*PED	Apr 30, 2029				
<u>MIRABEGRON - MYRBETRIQ GRANULES</u>						
N 213801 001	10058536	Mar 31, 2036	DP U-3108			
	10058536*PED	Sep 30, 2036				
<u>MITAPIVAT SULFATE - PYRUKYND</u>						
N 216196 001	10632114	May 03, 2032	U-3320		NCE	Feb 17, 2027
	11234976	Oct 11, 2038	U-3321		ODE-392	Feb 17, 2029
	11254652	Nov 21, 2038	DS DP			
	11793806	Apr 12, 2033	U-3320			
	11878049	Jul 31, 2041	U-3782			
	9193701	Oct 26, 2032	U-3319			
	9682080	May 03, 2032	U-3319			
	9980961	May 03, 2032	U-3319			
	RE49582	Feb 24, 2031	DS DP			
<u>MITAPIVAT SULFATE - PYRUKYND</u>						
N 216196 002	10632114	May 03, 2032	U-3320		NCE	Feb 17, 2027
	11234976	Oct 11, 2038	U-3321		ODE-392	Feb 17, 2029
	11254652	Nov 21, 2038	DS DP			
	11793806	Apr 12, 2033	U-3320			
	11878049	Jul 31, 2041	U-3782			
	9193701	Oct 26, 2032	U-3319			
	9682080	May 03, 2032	U-3319			
	9980961	May 03, 2032	U-3319			
	RE49582	Feb 24, 2031	DS DP			
<u>MITAPIVAT SULFATE - PYRUKYND</u>						
N 216196 003	10632114	May 03, 2032	U-3320		NCE	Feb 17, 2027
	11234976	Oct 11, 2038	U-3321		ODE-392	Feb 17, 2029
	11254652	Nov 21, 2038	DS DP			
	11793806	Apr 12, 2033	U-3320			
	11878049	Jul 31, 2041	U-3782			
	9193701	Oct 26, 2032	U-3319			
	9682080	May 03, 2032	U-3319			
	9980961	May 03, 2032	U-3319			
	RE49582	Feb 24, 2031	DS DP			
<u>MITOMYCIN - MITOSOL</u>						
N 022572 001	7806265	Feb 01, 2029	DP			
	8186511	Jul 19, 2026	DP			
	9205075	Jul 19, 2026	DP			
	9539241	Jan 02, 2028	DS DP U-2095			
	9649428	May 21, 2029	U-2095			
<u>MITOMYCIN - JELMYTO</u>						
N 211728 001	9040074	Jan 20, 2031	DP		ODE-289	Apr 15, 2027
	9950069	Jan 20, 2031	DP			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>MOBOCERTINIB SUCCINATE - EXKIVITY</u>						
N 215310	001	10227342	May 13, 2035	DS DP U-3220	NCE	Sep 15, 2026
		9796712	May 13, 2035	DS DP	ODE-374	Sep 15, 2028
<u>MOMELOTINIB DIHYDROCHLORIDE - OJJAARA</u>						
N 216873	001	11963962	Dec 02, 2040	U-3928	NCE	Sep 15, 2028
		8486941	Jan 03, 2030	DS DP U-1201	ODE-441	Sep 15, 2030
		9809559	Jun 11, 2035	U-1201		
		RE48285	Jun 11, 2035	DS DP U-1201		
<u>MOMELOTINIB DIHYDROCHLORIDE - OJJAARA</u>						
N 216873	002	11963962	Dec 02, 2040	U-3928	NCE	Sep 15, 2028
		8486941	Jan 03, 2030	DS DP U-1201	ODE-441	Sep 15, 2030
		9809559	Jun 11, 2035	U-1201		
		RE48285	Jun 11, 2035	DS DP U-1201		
<u>MOMELOTINIB DIHYDROCHLORIDE - OJJAARA</u>						
N 216873	003	11963962	Dec 02, 2040	U-3928	NCE	Sep 15, 2028
		8486941	Jan 03, 2030	DS DP U-1201	ODE-441	Sep 15, 2030
		9809559	Jun 11, 2035	U-1201		
		RE48285	Jun 11, 2035	DS DP U-1201		
<u>MOMETASONE FUROATE - SINUVA</u>						
N 209310	001	10232152	Nov 24, 2034	DP U-2272		
		10357640	Oct 03, 2031	U-2272		
		10406332	Mar 13, 2034	DP		
		7544192	Nov 29, 2026	U-2272		
		8025635	Jun 12, 2027	DP U-2272		
		8763222	Feb 08, 2032	DP		
		9585681	Apr 04, 2026	U-2272		
<u>MOMETASONE FUROATE; OLOPATADINE HYDROCHLORIDE - RYALTRIS</u>						
N 211746	001	10016443	Sep 04, 2034	U-3297	NP	Jan 13, 2025
		10376526	Sep 04, 2034	DP		
		10517880	Sep 04, 2034	DP U-3297		
		10548907	Sep 04, 2034	U-3297		
		10561672	Sep 04, 2034	DP		
		10646500	Sep 04, 2034	U-3297		
		10758550	Sep 04, 2034	U-3296		
		10765686	Sep 04, 2034	U-3295		
		11400101	Sep 04, 2034	U-3297		
		11679210	Sep 03, 2038	DP		
		12064442	Sep 04, 2034	U-3297		
		9078923	Sep 04, 2034	DP U-3297		
		9370483	Sep 04, 2034	DP		
		9750754	Sep 04, 2034	DP U-3297		
		9937189	Sep 04, 2034	DP		
<u>MONOMETHYL FUMARATE - BAFIERTAM</u>						
N 210296	001	10098863	Feb 27, 2035	DP U-1384		
		10105335	Feb 27, 2035	DP		
		10105336	Feb 27, 2035	DP U-1384		
		10105337	Feb 27, 2035	DP		
		10918615	Aug 12, 2035	DP U-1384		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>MONOMETHYL FUMARATE - BAFIERTAM</u>						
N 210296	001	10918616	Jun 03, 2035		U-1384	
		10918617	Aug 10, 2035		DP	
		10945985	Aug 14, 2035		DP	
		11590095	Mar 18, 2036		U-1384	
		11903918	Feb 27, 2035		U-1384	
		9326947	Feb 27, 2035		DP	
		9326965	Feb 27, 2035		U-1384	
		9511043	Feb 27, 2035		U-1384	
		9517209	Feb 27, 2035		DP	
		9566259	Feb 27, 2035		DP	
		9636318	Feb 27, 2035		U-1384	
		9636319	Feb 27, 2035		DP	
		9814691	Feb 27, 2035		U-1384	
		9814692	Feb 27, 2035		U-1384	
		9820960	Feb 27, 2035		DP	
		9820961	Feb 27, 2035		DP U-1384	
<u>MORPHINE SULFATE - MORPHINE SULFATE</u>						
N 204223	001	9072781	Mar 12, 2034		DP	
		9192608	Mar 12, 2034		U-43	
		9192608	Mar 12, 2034		U-55	
		9248229	Mar 12, 2034		DP	
<u>MORPHINE SULFATE - MORPHINE SULFATE</u>						
N 204223	002	9072781	Mar 12, 2034		DP	
		9192608	Mar 12, 2034		U-43	
		9192608	Mar 12, 2034		U-55	
		9248229	Mar 12, 2034		DP	
<u>MORPHINE SULFATE - MORPHINE SULFATE</u>						
N 204223	003	9072781	Mar 12, 2034		DP	
		9192608	Mar 12, 2034		U-43	
		9192608	Mar 12, 2034		U-55	
		9248229	Mar 12, 2034		DP	
<u>MORPHINE SULFATE - MORPHINE SULFATE</u>						
N 204223	004	9072781	Mar 12, 2034		DP	
		9192608	Mar 12, 2034		U-43	
		9192608	Mar 12, 2034		U-55	
		9248229	Mar 12, 2034		DP	
<u>MORPHINE SULFATE - MORPHINE SULFATE</u>						
N 204223	005	9072781	Mar 12, 2034		DP	
		9192608	Mar 12, 2034		U-43	
		9192608	Mar 12, 2034		U-55	
		9248229	Mar 12, 2034		DP	
<u>MORPHINE SULFATE - MORPHABOND ER</u>						
N 206544	001	10314788	Aug 12, 2028		DP	
		7955619	Aug 12, 2028		DP	
<u>MORPHINE SULFATE - MORPHABOND ER</u>						
N 206544	002	10314788	Aug 12, 2028		DP	
		7955619	Aug 12, 2028		DP	

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>MORPHINE SULFATE - MORPHABOND ER</u>						
N 206544	002	10314788	Aug 12, 2028	DP		
		7955619	Aug 12, 2028	DP		
<u>MORPHINE SULFATE - MORPHABOND ER</u>						
N 206544	003	10314788	Aug 12, 2028	DP		
		7955619	Aug 12, 2028	DP		
<u>MORPHINE SULFATE - MORPHABOND ER</u>						
N 206544	004	10314788	Aug 12, 2028	DP		
		7955619	Aug 12, 2028	DP		
<u>MORPHINE SULFATE - ARYMO ER</u>						
N 208603	001	9044402	Jul 01, 2033	DP U-1556		
		9549899	Jul 01, 2033	DP U-1556		
<u>MORPHINE SULFATE - ARYMO ER</u>						
N 208603	002	9044402	Jul 01, 2033	DP U-1556		
		9549899	Jul 01, 2033	DP U-1556		
<u>MORPHINE SULFATE - ARYMO ER</u>						
N 208603	003	9044402	Jul 01, 2033	DP U-1556		
		9549899	Jul 01, 2033	DP U-1556		
<u>MORPHINE SULFATE; NALTREXONE HYDROCHLORIDE - EMBEDA</u>						
N 022321	001	7682633	Jun 19, 2027	U-1510		
		7682634	Jun 19, 2027	DP		
		7815934	Dec 12, 2027	DP		
		8158156	Jun 19, 2027	U-1510		
		8623418	Nov 07, 2029	U-1640		
		8685443	Jul 03, 2025	U-1508		
		8685444	Jul 03, 2025	DP		
		8846104	Jun 19, 2027	DP		
		8877247	Jun 19, 2027	DP		
<u>MORPHINE SULFATE; NALTREXONE HYDROCHLORIDE - EMBEDA</u>						
N 022321	002	7682633	Jun 19, 2027	U-1510		
		7682634	Jun 19, 2027	DP		
		7815934	Dec 12, 2027	DP		
		8158156	Jun 19, 2027	U-1510		
		8623418	Nov 07, 2029	U-1640		
		8685443	Jul 03, 2025	U-1508		
		8685444	Jul 03, 2025	DP		
		8846104	Jun 19, 2027	DP		
		8877247	Jun 19, 2027	DP		
<u>MORPHINE SULFATE; NALTREXONE HYDROCHLORIDE - EMBEDA</u>						
N 022321	003	7682633	Jun 19, 2027	U-1510		
		7682634	Jun 19, 2027	DP		
		7815934	Dec 12, 2027	DP		
		8158156	Jun 19, 2027	U-1510		
		8623418	Nov 07, 2029	U-1640		
		8685443	Jul 03, 2025	U-1508		
		8685444	Jul 03, 2025	DP		
		8846104	Jun 19, 2027	DP		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>MORPHINE SULFATE; NALTREXONE HYDROCHLORIDE - EMBEDA</u>						
N 022321	003	8877247	Jun 19, 2027	DP		
<u>MORPHINE SULFATE; NALTREXONE HYDROCHLORIDE - EMBEDA</u>						
N 022321	004	7682633	Jun 19, 2027	U-1510		
		7682634	Jun 19, 2027	DP		
		7815934	Dec 12, 2027	DP		
		8158156	Jun 19, 2027	U-1510		
		8623418	Nov 07, 2029	U-1640		
		8685443	Jul 03, 2025	U-1508		
		8685444	Jul 03, 2025	DP		
		8846104	Jun 19, 2027	DP		
		8877247	Jun 19, 2027	DP		
<u>MORPHINE SULFATE; NALTREXONE HYDROCHLORIDE - EMBEDA</u>						
N 022321	005	7682633	Jun 19, 2027	U-1510		
		7682634	Jun 19, 2027	DP		
		7815934	Dec 12, 2027	DP		
		8158156	Jun 19, 2027	U-1510		
		8623418	Nov 07, 2029	U-1640		
		8685443	Jul 03, 2025	U-1508		
		8685444	Jul 03, 2025	DP		
		8846104	Jun 19, 2027	DP		
		8877247	Jun 19, 2027	DP		
<u>MORPHINE SULFATE; NALTREXONE HYDROCHLORIDE - EMBEDA</u>						
N 022321	006	7682633	Jun 19, 2027	U-1510		
		7682634	Jun 19, 2027	DP		
		7815934	Dec 12, 2027	DP		
		8158156	Jun 19, 2027	U-1510		
		8623418	Nov 07, 2029	U-1640		
		8685443	Jul 03, 2025	U-1508		
		8685444	Jul 03, 2025	DP		
		8846104	Jun 19, 2027	DP		
		8877247	Jun 19, 2027	DP		
<u>MOTIXAFORTIDE ACETATE - APHEXDA</u>						
N 217159	001				NCE	Sep 08, 2028
					ODE-442	Sep 08, 2030
<u>MOXIDECTIN - MOXIDECTIN</u>						
N 210867	001				ODE-193	Jun 13, 2025
<u>MOXIFLOXACIN HYDROCHLORIDE - MOXEZA</u>						
N 022428	001	8450311	May 29, 2029	DP		
		9114168	May 29, 2029	DP		
<u>MYCOPHENOLATE MOFETIL - MYHIBBIN</u>						
N 216482	001	11931455	Aug 16, 2039	DP	U-1752	
		12097284	Aug 16, 2039	DP		
		12097285	Aug 16, 2039	DP	U-1752	
<u>NAFTIFINE HYDROCHLORIDE - NAFTIN</u>						
N 204286	001	10166205	Jan 31, 2033	DP		
		10166206	Jan 31, 2033	DP		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>NAFTIFINE HYDROCHLORIDE - NAFTIN</u>						
N 204286	001 10695303	Jan 31, 2033	DP			
	10729667	Jan 31, 2033	DP			
	8778365	Jan 31, 2033	DP			
	9161914	Jan 31, 2033		U-540		
<u>NALBUPHINE HYDROCHLORIDE - NALBUPHINE HYDROCHLORIDE</u>						
A 216049	001				CGT	Mar 30, 2025
<u>NALBUPHINE HYDROCHLORIDE - NALBUPHINE HYDROCHLORIDE</u>						
A 216049	002				CGT	Mar 30, 2025
<u>NALBUPHINE HYDROCHLORIDE - NALBUPHINE HYDROCHLORIDE</u>						
A 216050	001				CGT	Mar 24, 2025
<u>NALBUPHINE HYDROCHLORIDE - NALBUPHINE HYDROCHLORIDE</u>						
A 216050	002				CGT	Mar 24, 2025
<u>NALDEMEDINE TOSYLATE - SYMPROIC</u>						
N 208854	001 10952968	May 13, 2033	DS DP			
	9108975	Nov 11, 2031	DS DP			
	RE46365	Jan 11, 2028	DS DP			
	RE46375	Oct 05, 2026	DS DP	U-1185		
<u>NALMEFENE HYDROCHLORIDE - OPVEE</u>						
N 217470	001 11458091	Jul 10, 2038	DP	U-3630	NP	May 22, 2026
<u>NALMEFENE HYDROCHLORIDE - ZURNAL (AUTOINJECTOR)</u>						
N 218590	001 10279131	Jul 31, 2031	DP		NP	Aug 07, 2027
	10357609	Aug 21, 2031	DP			
	10478560	Jan 24, 2026	DP			
	10881798	Feb 11, 2034	DP			
	10905827	Aug 21, 2031	DP			
	11185642	Aug 28, 2031	DP			
	11191908	Oct 18, 2035	DP			
	11446440	Aug 21, 2031	DP			
	11446441	Jan 24, 2026	DP			
	11813435	Feb 25, 2035	DP			
	11857547	Nov 05, 2039	DP			
	11865112	Nov 05, 2039	DP	U-3630		
	8021335	Oct 04, 2026	DP			
	8496619	Aug 21, 2031	DP			
	8562564	Jan 24, 2026	DP			
	9180259	Jan 24, 2026	DP			
	9364610	Aug 21, 2031	DP			
	9364611	Aug 21, 2031	DP			
	9446195	Aug 21, 2031	DP			
	9533102	Jan 24, 2026	DP			
	9629959	Jan 24, 2026	DP			
<u>NALOXEGOL OXALATE - MOVANTIK</u>						
N 204760	001 7786133	Sep 16, 2028	DS DP			
	9012469	Apr 02, 2032	DS DP			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>NALOXEGOL OXALATE - MOVANTIK</u>						
N 204760	002 7786133	Sep 16, 2028	DS DP			
	9012469	Apr 02, 2032	DS DP			
<u>NALOXONE HYDROCHLORIDE - EVZIO</u>						
N 205787	001 10143972	May 24, 2031		U-2476		
	10220158	Mar 20, 2035	DP	U-2500		
	10322239	Feb 28, 2031		U-1907		
	10335549	Apr 30, 2025	DP			
	10960155	Jun 25, 2026	DP			
	11590286	Dec 12, 2026	DP			
	7731686	Jun 10, 2026	DP			
	7731690	Jan 15, 2025	DP			
	7749194	Oct 30, 2028	DP			
	7947017	Mar 12, 2028	DP			
	8016788	Mar 21, 2025	DP			
	8021344	Nov 02, 2029	DP			
	8206360	Feb 27, 2027	DP			
	8226610	Apr 10, 2029	DP			
	8231573	Nov 25, 2028	DP			
	8627816	Feb 04, 2032	DP			
	8926594	Mar 31, 2026	DP			
	8939943	Feb 28, 2031	DP			
	9022022	Feb 28, 2031	DP			
	9238108	Feb 20, 2027	DP			
	9278182	Feb 01, 2026	DP			
	9474869	Feb 28, 2031	DP	U-1907		
	9517307	Jul 18, 2034	DP	U-1925		
	9724471	May 23, 2027	DP	U-2092		
<u>NALOXONE HYDROCHLORIDE - NARCAN</u>						
N 208411	002 9480644	Mar 16, 2035	DP	U-1903		Y
	9707226	Mar 16, 2035	DP	U-1903		Y
<u>NALOXONE HYDROCHLORIDE - REXTOVY</u>						
N 208969	001				NP	Mar 07, 2026
<u>NALOXONE HYDROCHLORIDE - EVZIO (AUTOINJECTOR)</u>						
N 209862	001 10143792	May 24, 2031		U-2476		
	10220158	Mar 20, 2035	DP	U-2500		
	10322239	Feb 28, 2031		U-1907		
	10335549	Apr 30, 2025	DP			
	10960155	Jun 25, 2026	DP			
	11590286	Dec 12, 2026	DP			
	7731686	Jun 01, 2026	DP			
	7731690	Jan 15, 2025	DP			
	7749194	Oct 30, 2028	DP			
	7947017	Mar 12, 2028	DP			
	8016788	Mar 21, 2025	DP			
	8021344	Nov 02, 2029	DP			
	8206360	Feb 27, 2027	DP			
	8226610	Apr 10, 2029	DP			
	8231573	Nov 25, 2028	DP			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>NALOXONE HYDROCHLORIDE - EVZIO (AUTOINJECTOR)</u>						
N 209862	001 8627816	Feb 04, 2032	DP			
	8926594	Mar 31, 2026	DP			
	8939943	Feb 28, 2031	DP			
	9022022	Feb 28, 2031	DP			
	9238108	Feb 20, 2027	DP			
	9278182	Feb 01, 2026	DP			
	9474869	Feb 28, 2031	DP U-1907			
	9517307	Jul 18, 2034	DP U-1925			
	9724471	May 23, 2027	DP U-2092			
<u>NALOXONE HYDROCHLORIDE - KLOXXADO</u>						
N 212045	001 10722510	Aug 26, 2034	DP U-3110			
	10973814	Aug 26, 2034	DP U-3110			
	11135155	Aug 26, 2034	DP			
	11617713	Aug 26, 2034	DP U-3110			
	11628139	Aug 26, 2034	DP U-3110			
	11975096	Aug 26, 2034	DP			
<u>NALOXONE HYDROCHLORIDE - ZIMHI</u>						
N 212854	001 11027072	May 24, 2039	DP			
	11571518	Jun 14, 2041	DP U-3515			
	11571518	Jun 14, 2041	DP U-3516			
	11571518	Jun 14, 2041	DP U-3517			
<u>NALOXONE HYDROCHLORIDE - NALOXONE HYDROCHLORIDE (AUTOINJECTOR)</u>						
N 215457	001 10143792	May 24, 2031	U-2476			
	10322239	Feb 28, 2031	U-1907			
	10335549	Apr 30, 2025	DP			
	11590286	Dec 12, 2026	DP			
	7731690	Jan 15, 2025	DP			
	7947017	Mar 12, 2028	DP			
	8016788	Mar 21, 2025	DP			
	8939943	Feb 28, 2031	DP			
	9022022	Feb 28, 2031	DP			
	9474869	Feb 28, 2031	DP U-1907			
	9814838	Feb 28, 2031	DP			
<u>NALOXONE HYDROCHLORIDE - REZENOPY</u>						
N 215487	001				NP	Apr 19, 2027
<u>NALOXONE HYDROCHLORIDE - RIVIVE</u>						
N 217722	001 11020343	May 11, 2032	U-3671			
	11806428	May 11, 2032	DP			
<u>NALOXONE HYDROCHLORIDE; OXYCODONE HYDROCHLORIDE - TARGINIQ</u>						
N 205777	001 9073933	Mar 30, 2025	DS			
	9522919	Mar 30, 2025	DS DP			
<u>NALOXONE HYDROCHLORIDE; OXYCODONE HYDROCHLORIDE - TARGINIQ</u>						
N 205777	002 9073933	Mar 30, 2025	DS			
	9522919	Mar 30, 2025	DS DP			
<u>NALOXONE HYDROCHLORIDE; OXYCODONE HYDROCHLORIDE - TARGINIQ</u>						
N 205777	003 9073933	Mar 30, 2025	DS			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>NALOXONE HYDROCHLORIDE; OXYCODONE HYDROCHLORIDE - TARGINIQ</u>						
N 205777	003 9522919	Mar 30, 2025	DS DP			
<u>NALTREXONE - VIVITROL</u>						
N 021897	001 7919499	Oct 15, 2029	U-1123			
	7919499	Oct 15, 2029	U-1124			
<u>NALTREXONE HYDROCHLORIDE; OXYCODONE HYDROCHLORIDE - TROXYCA ER</u>						
N 207621	001 7815934	Dec 12, 2027	DP			
	8685443	Jul 03, 2025	U-1508			
<u>NALTREXONE HYDROCHLORIDE; OXYCODONE HYDROCHLORIDE - TROXYCA ER</u>						
N 207621	002 7815934	Dec 12, 2027	DP			
	8685443	Jul 03, 2025	U-1508			
<u>NALTREXONE HYDROCHLORIDE; OXYCODONE HYDROCHLORIDE - TROXYCA ER</u>						
N 207621	003 7815934	Dec 12, 2027	DP			
	8685443	Jul 03, 2025	U-1508			
<u>NALTREXONE HYDROCHLORIDE; OXYCODONE HYDROCHLORIDE - TROXYCA ER</u>						
N 207621	004 7815934	Dec 12, 2027	DP			
	8685443	Jul 03, 2025	U-1508			
<u>NALTREXONE HYDROCHLORIDE; OXYCODONE HYDROCHLORIDE - TROXYCA ER</u>						
N 207621	005 7815934	Dec 12, 2027	DP			
	8685443	Jul 03, 2025	U-1508			
<u>NALTREXONE HYDROCHLORIDE; OXYCODONE HYDROCHLORIDE - TROXYCA ER</u>						
N 207621	006 7815934	Dec 12, 2027	DP			
	8685443	Jul 03, 2025	U-1508			
<u>NAPROXEN SODIUM - NAPROXEN SODIUM</u>						
N 021920	001 10022344	Mar 03, 2026	DP U-1731			
	10022344	Mar 03, 2026	DP U-1732			
	10028925	Mar 03, 2026	DP U-1731			
	10028925	Mar 03, 2026	DP U-1732			
	11090280	Mar 03, 2026	DP U-1731			
	11090280	Mar 03, 2026	DP U-1732			
	9693978	Mar 03, 2026	DP			
	9693979	Mar 03, 2026	DP			
<u>NAPROXEN SODIUM; SUMATRIPTAN SUCCINATE - TREXIMET</u>						
N 021926	001 7332183	Oct 02, 2025	DP U-867			
	7332183*PED	Apr 02, 2026				
<u>NAPROXEN SODIUM; SUMATRIPTAN SUCCINATE - TREXIMET</u>						
N 021926	002 7332183	Oct 02, 2025	DP U-1719			
	7332183*PED	Apr 02, 2026				
<u>NEBIVOLOL HYDROCHLORIDE; VALSARTAN - BYVALSON</u>						
N 206302	001 7803838	Aug 29, 2026	DP			
	7838552	Oct 04, 2027	U-185			
<u>NEDOSIRAN SODIUM - RIVFLOZA</u>						
N 215842	001 10351854	Oct 09, 2035	DS DP		NCE	Sep 29, 2028
	10738311	Oct 09, 2035	DS DP U-3709		ODE-443	Sep 29, 2030
	11053502	Oct 29, 2035	DS DP			
	11286488	Oct 12, 2038	DS DP U-3709			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>NEDOSIRAN SODIUM - RIVFLOZA</u>						
N 215842	001	11359203	Oct 09, 2035	DS DP U-3709		
		11661604	Oct 12, 2038	DS DP U-3709		
<u>NEDOSIRAN SODIUM - RIVFLOZA</u>						
N 215842	002	10351854	Oct 09, 2035	DS DP	NCE	Sep 29, 2028
		10738311	Oct 09, 2035	DS DP U-3709	ODE-443	Sep 29, 2030
		11053502	Oct 29, 2035	DS DP		
		11286488	Oct 12, 2038	DS DP U-3709		
		11359203	Oct 09, 2035	DS DP U-3709		
		11661604	Oct 12, 2038	DS DP U-3709		
<u>NEDOSIRAN SODIUM - RIVFLOZA</u>						
N 215842	003	10351854	Oct 09, 2035	DS DP	NCE	Sep 29, 2028
		10738311	Oct 09, 2035	DS DP U-3709	ODE-443	Sep 29, 2030
		11053502	Oct 29, 2035	DS DP		
		11286488	Oct 12, 2038	DS DP U-3709		
		11359203	Oct 09, 2035	DS DP U-3709		
		11661604	Oct 12, 2038	DS DP U-3709		
<u>NEPAFENAC - NEVANAC</u>						
N 021862	001	7834059	Jan 31, 2027	U-1095		
		8071648	Dec 02, 2025	DP		
		8324281	Dec 02, 2025	DP		
<u>NEPAFENAC - ILEVRO</u>						
N 203491	001	8921337	Mar 31, 2032	DP		
		9662398	Dec 01, 2030	DP		
<u>NERATINIB MALEATE - NERLYNX</u>						
N 208051	001	10035788	Oct 15, 2028	U-2043		
		10035788	Oct 15, 2028	U-3047		
		10035788	Oct 15, 2028	U-3097		
		7399865	Dec 29, 2030	DS DP		
		7982043	Oct 08, 2025	U-2043		
		7982043	Oct 08, 2025	U-3047		
		7982043	Oct 08, 2025	U-3097		
		8518446	Nov 20, 2030	DP U-2043		
		8518446	Nov 20, 2030	DP U-3047		
		8518446	Nov 20, 2030	DP U-3097		
		8669273	Jul 18, 2031	U-3047		
		8790708	Nov 05, 2030	DP U-2043		
		8790708	Nov 05, 2030	DP U-3047		
		8790708	Nov 05, 2030	DP U-3097		
		9139558	Oct 15, 2028	U-2043		
		9139558	Oct 15, 2028	U-3047		
		9139558	Oct 15, 2028	U-3097		
		9211291	Mar 24, 2030	U-2043		
		9211291	Mar 24, 2030	U-3097		
		9265784	Aug 04, 2029	U-3047		
		9630946	Oct 15, 2028	U-2043		
		9630946	Oct 15, 2028	U-3047		
		9630946	Oct 15, 2028	U-3097		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>NETARSUDIL MESYLATE - RHOPRESSA</u>						
N 208254	001	10174017	Jan 27, 2030	DS DP	U-1524	
		10532993	Jul 11, 2026		U-1524	
		10588901	Mar 14, 2034	DS DP	U-1524	
		10654844	Jan 27, 2030	DS DP	U-1524	
		10882840	Jul 11, 2026		U-1524	
		11020385	Mar 14, 2034		U-1524	
		11021456	Jul 11, 2026		U-1524	
		11028081	Jan 27, 2030		U-1524	
		11185538	Mar 14, 2034	DP		
		11618748	Jan 27, 2030		U-1524	
		8394826	Nov 10, 2030	DS DP	U-1524	
		8450344	Jul 11, 2026	DS DP	U-1524	
		9096569	Jul 11, 2026	DS DP	U-1524	
		9415043	Mar 14, 2034	DS		
		9931336	Mar 14, 2034	DS DP	U-1524	
<u>NETUPITANT; PALONOSETRON HYDROCHLORIDE - AKYNZEO</u>						
N 205718	001	10233154	Sep 25, 2035	DS		
		10676440	Sep 25, 2035	DS DP		
		10828297	Dec 17, 2030		U-2293	
		10961195	Sep 25, 2035	DS DP		
		11559523	Nov 18, 2030	DP	U-3522	
		12042494	Nov 18, 2030	DP	U-3522	
		8623826	Nov 18, 2030		U-2293	
		8951969	Nov 18, 2030	DP		
		9186357	Nov 18, 2030		U-2293	
		9271975	Sep 09, 2031		U-2293	
		9943515	Nov 18, 2030		U-2293	
		9951016	Sep 25, 2035	DS DP		
<u>NEVIRAPINE - VIRAMUNE XR</u>						
N 201152	001	8460704	Mar 12, 2029		U-1409	
<u>NICARDIPINE HYDROCHLORIDE - NICARDIPINE HYDROCHLORIDE IN 0.9% SODIUM CHLORIDE</u>						
A 215592	001				CGT	May 21, 2025
<u>NICARDIPINE HYDROCHLORIDE - NICARDIPINE HYDROCHLORIDE IN 0.9% SODIUM CHLORIDE</u>						
A 215592	002				CGT	May 21, 2025
<u>NICARDIPINE HYDROCHLORIDE - CARDENE IN 4.8% DEXTROSE IN PLASTIC CONTAINER</u>						
N 019734	002	10758616	Apr 18, 2027	DP		
		11547758	Apr 18, 2027		U-1029	
		7612102	Dec 26, 2027	DP		
		7659291	Apr 18, 2027		U-1029	
		8455524	Apr 18, 2027		U-1029	
		9364564	Dec 26, 2027	DP		
<u>NICARDIPINE HYDROCHLORIDE - CARDENE IN 0.86% SODIUM CHLORIDE IN PLASTIC CONTAINER</u>						
N 019734	003	10758616	Apr 18, 2027	DP		
		11547758	Apr 18, 2027		U-1029	
		7612102	Dec 26, 2027	DP		
		7659291	Apr 18, 2027		U-1029	
		8455524	Apr 18, 2027		U-1029	

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>NICARDIPINE HYDROCHLORIDE - CARDENE IN 0.86% SODIUM CHLORIDE IN PLASTIC CONTAINER</u>						
N 019734	003	9364564	Dec 26, 2027	DP		
<u>NICARDIPINE HYDROCHLORIDE - CARDENE IN 0.83% SODIUM CHLORIDE IN PLASTIC CONTAINER</u>						
N 019734	004	10758616	Apr 18, 2027	DP		
		11547758	Apr 18, 2027		U-1029	
		7612102	Dec 26, 2027	DP		
		7659291	Apr 18, 2027		U-1029	
		8455524	Apr 18, 2027		U-1029	
		9364564	Dec 26, 2027	DP		
<u>NICARDIPINE HYDROCHLORIDE - CARDENE IN 5.0% DEXTROSE IN PLASTIC CONTAINER</u>						
N 019734	005	7612102	Dec 26, 2027	DP		
		7659291	Apr 18, 2027		U-1029	
		8455524	Apr 18, 2027		U-1029	
		9364564	Dec 26, 2027	DP		
<u>NICOTINE POLACRILEX - NICORETTE</u>						
N 018612	002	8323683	Apr 30, 2028			
<u>NICOTINE POLACRILEX - NICORETTE</u>						
N 020066	002	8323683	Apr 30, 2028	DP		
<u>NICOTINE POLACRILEX - NICORETTE</u>						
N 022360	001	8501164	Jun 14, 2029	DP		
		8940772	Apr 30, 2029	DP		
<u>NICOTINE POLACRILEX - NICORETTE</u>						
N 022360	002	8501164	Jun 14, 2029	DP		
		8940772	Apr 30, 2029	DP		
<u>NIFURTIMOX - LAMPIT</u>						
N 213464	001				NCE	Aug 06, 2025
					ODE-319	Aug 06, 2027
<u>NIFURTIMOX - LAMPIT</u>						
N 213464	002				NCE	Aug 06, 2025
					ODE-319	Aug 06, 2027
<u>NILOTINIB HYDROCHLORIDE - TASIGNA</u>						
N 022068	001	8163904	Aug 23, 2028	DS DP	ODE-171	Mar 22, 2025
		8163904*PED	Feb 23, 2029		ODE-172	Mar 22, 2025
		8293756	Sep 25, 2027	DP	ODE-380	Sep 23, 2028
		8293756*PED	Mar 25, 2028		PED	Sep 22, 2025
		8389537	Jul 18, 2026	U-1374	PED	Sep 22, 2025
		8389537*PED	Jan 18, 2027		PED	Mar 23, 2029
		8415363	Jul 18, 2026	DS DP	U-1374	
		8415363	Jul 18, 2026	DS DP	U-1407	
		8415363*PED	Jan 18, 2027			
		8501760	Jul 18, 2026	DP		
		8501760*PED	Jan 18, 2027			
		9061029	Apr 07, 2032	DS	U-1374	
		9061029	Apr 07, 2032	DS	U-3231	
		9061029*PED	Oct 07, 2032			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>NILOTINIB HYDROCHLORIDE - TASIGNA</u>						
N 022068	002	8163904	Aug 23, 2028	DS DP	ODE-171	Mar 22, 2025
		8163904*PED	Feb 23, 2029		ODE-172	Mar 22, 2025
		8293756	Sep 25, 2027	DP	ODE-380	Sep 23, 2028
		8293756*PED	Mar 25, 2028		PED	Sep 22, 2025
		8389537	Jul 18, 2026	U-1374	PED	Sep 22, 2025
		8389537*PED	Jan 18, 2027		PED	Mar 23, 2029
		8415363	Jul 18, 2026	DS DP U-1374		
		8415363	Jul 18, 2026	DS DP U-1407		
		8415363*PED	Jan 18, 2027			
		8501760	Jul 18, 2026	DP		
		8501760*PED	Jan 18, 2027			
		9061029	Apr 07, 2032	DS U-1374		
		9061029	Apr 07, 2032	DS U-3231		
		9061029*PED	Oct 07, 2032			
<u>NILOTINIB HYDROCHLORIDE - TASIGNA</u>						
N 022068	003	8163904	Aug 23, 2028	DS DP	ODE-171	Mar 22, 2025
		8163904*PED	Feb 23, 2029		ODE-172	Mar 22, 2025
		8293756	Sep 25, 2027	DP	ODE-380	Sep 23, 2028
		8293756*PED	Mar 25, 2028		PED	Sep 22, 2025
		8389537	Jul 18, 2026	U-1374	PED	Sep 22, 2025
		8389537*PED	Jan 18, 2027		PED	Mar 23, 2029
		8415363	Jul 18, 2026	DS DP U-1374		
		8415363	Jul 18, 2026	DS DP U-1407		
		8415363*PED	Jan 18, 2027			
		8501760	Jul 18, 2026	DP		
		8501760*PED	Jan 18, 2027			
		9061029	Apr 07, 2032	DS U-1374		
		9061029	Apr 07, 2032	DS U-3231		
		9061029*PED	Oct 07, 2032			
<u>NILOTINIB TARTRATE - DANZITEN</u>						
N 219293	001	10874671	Feb 18, 2040	DP		
		11793809	Feb 18, 2040	DP		
<u>NILOTINIB TARTRATE - DANZITEN</u>						
N 219293	002	10874671	Feb 18, 2040	DP		
		11793809	Feb 18, 2040	DP		
<u>NIMODIPINE - NYMALIZE</u>						
N 203340	002	10342787	Apr 16, 2038	DP U-2804		
		10576070	Apr 16, 2038	DP U-2804		
		11207306	Apr 16, 2038	U-2804		
		11413277	Apr 16, 2038	DP U-2804		
		11806338	Apr 16, 2038	DP		
<u>NINTEDANIB ESYLATE - OFEV</u>						
N 205832	001	10105323	Jun 04, 2029	DP	ODE-261	Sep 06, 2026
		10105323*PED	Dec 04, 2029		PED	Mar 06, 2027
		10154990	Jan 08, 2026	U-2620		
		10154990*PED	Jul 08, 2026			
		6762180	Oct 01, 2025	DS DP		
		6762180*PED	Apr 01, 2026			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>NINTEDANIB ESYLATE - OFEV</u>						
N 205832	001 9907756	Jun 07, 2029	DP			
	9907756*PED	Dec 07, 2029				
<u>NINTEDANIB ESYLATE - OFEV</u>						
N 205832	002 10105323	Jun 04, 2029	DP		ODE-261	Sep 06, 2026
	10105323*PED	Dec 04, 2029			PED	Mar 06, 2027
	10154990	Jan 08, 2026	U-2620			
	10154990*PED	Jul 08, 2026				
	6762180	Oct 01, 2025	DS DP			
	6762180*PED	Apr 01, 2026				
	9907756	Jun 07, 2029	DP			
	9907756*PED	Dec 07, 2029				
<u>NIRAPARIB TOSYLATE - ZEJULA</u>						
N 208447	001 11091459	Mar 27, 2038	DP		ODE-277	Oct 23, 2026
	11673877	Mar 27, 2038	DP U-3646		ODE-295	Apr 29, 2027
	11673877	Mar 27, 2038	DP U-3647			
	8071579	Aug 12, 2027	U-2655			
	8071623	Mar 27, 2031	DS DP			
	8143241	Aug 12, 2027	U-2655			
	8436185	Apr 24, 2029	DS			
	8859562	Aug 04, 2031	U-2655			
<u>NIRAPARIB TOSYLATE - ZEJULA</u>						
N 214876	001 11091459	Mar 27, 2038	DP			
	11673877	Mar 27, 2038	DP U-3646			
	11673877	Mar 27, 2038	DP U-3647			
	11730725	Jan 04, 2039	DP			
	8071579	Aug 12, 2027	U-2655			
	8071623	Mar 27, 2031	DS DP			
	8143241	Aug 12, 2027	U-2655			
	8436185	Apr 24, 2029	DS			
	8859562	Aug 04, 2031	U-2655			
<u>NIRAPARIB TOSYLATE - ZEJULA</u>						
N 214876	002 11091459	Mar 27, 2038	DP			
	11673877	Mar 27, 2038	DP U-3646			
	11673877	Mar 27, 2038	DP U-3647			
	11730725	Jan 04, 2039	DP			
	8071579	Aug 12, 2027	U-2655			
	8071623	Mar 27, 2031	DS DP			
	8143241	Aug 12, 2027	U-2655			
	8436185	Apr 24, 2029	DS			
	8859562	Aug 04, 2031	U-2655			
<u>NIRAPARIB TOSYLATE - ZEJULA</u>						
N 214876	003 11091459	Mar 27, 2038	DP			
	11673877	Mar 27, 2038	DP U-3646			
	11673877	Mar 27, 2038	DP U-3647			
	11730725	Jan 04, 2039	DP			
	8071579	Aug 12, 2027	U-2655			
	8071623	Mar 27, 2031	DS DP			
	8143241	Aug 12, 2027	U-2655			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>NIRAPARIB TOSYLATE - ZEJULA</u>						
N 214876	003 8436185	Apr 24, 2029	DS			
	8859562	Aug 04, 2031			U-2655	
<u>NIRMATRELVIR; RITONAVIR - PAXLOVID (COPACKAGED)</u>						
N 217188	001 11351149	Aug 05, 2041	DS DP	U-3629	NCE	May 25, 2028
	11541034	Oct 31, 2041		U-3629		
<u>NIROGACESTAT HYDROBROMIDE - OGSIVEO</u>						
N 217677	001 10590087	Aug 09, 2039	DS		NCE	Nov 27, 2028
	10710966	Aug 09, 2039	DS	U-3754	ODE-452	Nov 27, 2030
	10941118	Aug 09, 2039	DS	U-3754		
	11504354	Jul 08, 2042		DP		
	11612588	Jul 08, 2042		DP U-3754		
	11807611	Sep 08, 2042		DP U-3754		
	11820748	Aug 09, 2039		DP		
	11844780	Sep 08, 2042		DP U-3754		
	11845732	Aug 09, 2039	DS	U-3754		
	11872211	May 19, 2043		U-3754		
	11884634	Aug 09, 2039		DP		
	11884635	Aug 09, 2039		DP		
	11905255	Aug 09, 2039		DP		
	11925619	May 19, 2043		U-3754		
	11925620	May 19, 2043		U-3754		
	11938116	May 19, 2043		U-3754		
	11951096	May 19, 2043		U-3754		
	11957662	May 19, 2043		U-3754		
	12011434	May 19, 2043		U-3754		
	12011435	May 19, 2043		U-3754		
	12036207	May 19, 2043		U-3754		
	12110277	Jul 08, 2042		DP		
	12116347	Aug 09, 2039		DP		
	12138246	May 19, 2043		U-3754		
	7342118	Aug 18, 2025	DS			
	7795447	Aug 18, 2025	DS			
	7951958	Mar 11, 2025	DS			
<u>NIROGACESTAT HYDROBROMIDE - OGSIVEO</u>						
N 217677	002 10590087	Aug 09, 2039	DS		NCE	Nov 27, 2028
	10710966	Aug 09, 2039	DS	U-3754	ODE*	Nov 27, 2030
	10941118	Aug 09, 2039	DS	U-3754		
	11504354	Jul 08, 2042		DP		
	11612588	Jul 08, 2042		DP U-3754		
	11807611	Sep 08, 2042		DP U-3754		
	11820748	Aug 09, 2039		DP		
	11844780	Sep 08, 2042		DP U-3754		
	11845732	Aug 09, 2039	DS	U-3754		
	11872211	May 19, 2043		U-3754		
	11884634	Aug 09, 2039		DP		
	11884635	Aug 09, 2039		DP		
	11905255	Aug 09, 2039		DP		
	11925619	May 19, 2043		U-3754		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>NIROGACESTAT HYDROBROMIDE - OGSIVEO</u>						
N 217677	002	11925620	May 19, 2043		U-3754	
		11938116	May 19, 2043		U-3754	
		11951096	May 19, 2043		U-3754	
		11957662	May 19, 2043		U-3754	
		12011434	May 19, 2043		U-3754	
		12011435	May 19, 2043		U-3754	
		12036207	May 19, 2043		U-3754	
		12110277	Jul 08, 2042	DP		
		12116347	Aug 09, 2039	DP		
		12138246	May 19, 2043		U-3754	
		7342118	Aug 18, 2025	DS		
		7795447	Aug 18, 2025	DS		
		7951958	Mar 11, 2025	DS		
<u>NIROGACESTAT HYDROBROMIDE - OGSIVEO</u>						
N 217677	003	10590087	Aug 09, 2039	DS		NCE Nov 27, 2028
		10710966	Aug 09, 2039	DS	U-3754	ODE* Nov 27, 2030
		10941118	Aug 09, 2039	DS	U-3754	
		11504354	Jul 08, 2042	DP		
		11612588	Jul 08, 2042	DP	U-3754	
		11807611	Sep 08, 2042	DP	U-3754	
		11820748	Aug 09, 2039	DP		
		11844780	Sep 08, 2042	DP	U-3754	
		11845732	Aug 09, 2039	DS	U-3754	
		11872211	May 19, 2043		U-3754	
		11884634	Aug 09, 2039	DP		
		11884635	Aug 09, 2039	DP		
		11905255	Aug 09, 2039	DP		
		11925619	May 19, 2043		U-3754	
		11925620	May 19, 2043		U-3754	
		11938116	May 19, 2043		U-3754	
		11951096	May 19, 2043		U-3754	
		11957662	May 19, 2043		U-3754	
		12011434	May 19, 2043		U-3754	
		12011435	May 19, 2043		U-3754	
		12036207	May 19, 2043		U-3754	
		12110277	Jul 08, 2042	DP		
		12116347	Aug 09, 2039	DP		
		12138246	May 19, 2043		U-3754	
		7342118	Aug 18, 2025	DS		
		7795447	Aug 18, 2025	DS		
		7951958	Mar 11, 2025	DS		
<u>NITISINONE - ORFADIN</u>						
N 206356	001	9301932	Feb 28, 2033	DP	U-1836	
<u>NITISINONE - NITYR</u>						
N 209449	001	10328029	Jan 05, 2035	DP	U-1836	
<u>NITISINONE - NITYR</u>						
N 209449	002	10328029	Jan 05, 2035	DP	U-1836	

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>NITISINONE - NITYR</u>						
N 209449	003 10328029	Jan 05, 2035	DP U-1836			
<u>NITRIC OXIDE - INOMAX</u>						
N 020845	002 8282966	Jun 30, 2029	U-1286			
	8291904	Jan 06, 2031	DP U-1226			
	8293284	Jun 30, 2029	U-1286			
	8431163	Jun 30, 2029	U-1286			
	8431163*PED	Dec 30, 2029				
	8573209	Jan 06, 2031	DP			
	8573209*PED	Jul 06, 2031				
	8573210	Jan 06, 2031	DP U-1453			
	8573210*PED	Jul 06, 2031				
	8776794	Jan 06, 2031	DP U-1226			
	8776794*PED	Jul 06, 2031				
	8776795	Jan 06, 2031	DP U-1226			
	8776795*PED	Jul 06, 2031				
	8795741	Jun 30, 2029	U-1286			
	8795741*PED	Dec 30, 2029				
	8846112	Jun 30, 2029	U-1286			
	8846112*PED	Dec 30, 2029				
<u>NITRIC OXIDE - INOMAX</u>						
N 020845	003 11931377	Jun 03, 2029	DS DP U-3903			
	11931377*PED	Dec 03, 2029				
	8282966	Jun 30, 2029	U-1286			
	8291904	Jan 06, 2031	DP U-1226			
	8293284	Jun 30, 2029	U-1286			
	8431163	Jun 30, 2029	U-1286			
	8431163*PED	Dec 30, 2029				
	8573209	Jan 06, 2031	DP			
	8573209*PED	Jul 06, 2031				
	8573210	Jan 06, 2031	DP U-1453			
	8573210*PED	Jul 06, 2031				
	8776794	Jan 06, 2031	DP U-1226			
	8776794*PED	Jul 06, 2031				
	8776795	Jan 06, 2031	DP U-1226			
	8776795*PED	Jul 06, 2031				
	8795741	Jun 30, 2029	U-1286			
	8795741*PED	Dec 30, 2029				
	8846112	Jun 30, 2029	U-1286			
	8846112*PED	Dec 30, 2029				
	9265911	Jan 06, 2031	DP U-1824			
	9265911*PED	Jul 06, 2031				
	9279794	Feb 19, 2034	DP U-1823			
	9279794*PED	Aug 19, 2034				
	9295802	Jan 06, 2031	DP U-1226			
	9295802*PED	Jul 06, 2031				
	9408993	Jan 06, 2031	DP U-1824			
	9408993*PED	Jul 06, 2031				
	9770570	May 03, 2036	U-2148			
	9770570*PED	Nov 03, 2036				

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>NITRIC OXIDE - INOMAX</u>						
N 020845 003	11931377	Jun 03, 2029	DS DP U-3903			
	11931377*PED	Dec 03, 2029				
	8282966	Jun 30, 2029		U-1286		
	8291904	Jan 06, 2031	DP U-1226			
	8293284	Jun 30, 2029		U-1286		
	8431163	Jun 30, 2029		U-1286		
	8431163*PED	Dec 30, 2029				
	8573209	Jan 06, 2031	DP			
	8573209*PED	Jul 06, 2031				
	8573210	Jan 06, 2031	DP U-1453			
	8573210*PED	Jul 06, 2031				
	8776794	Jan 06, 2031	DP U-1226			
	8776794*PED	Jul 06, 2031				
	8776795	Jan 06, 2031	DP U-1226			
	8776795*PED	Jul 06, 2031				
	8795741	Jun 30, 2029		U-1286		
	8795741*PED	Dec 30, 2029				
	8846112	Jun 30, 2029		U-1286		
	8846112*PED	Dec 30, 2029				
	9265911	Jan 06, 2031	DP U-1824			
	9265911*PED	Jul 06, 2031				
	9279794	Feb 19, 2034	DP U-1823			
	9279794*PED	Aug 19, 2034				
	9295802	Jan 06, 2031	DP U-1226			
	9295802*PED	Jul 06, 2031				
	9408993	Jan 06, 2031	DP U-1824			
	9408993*PED	Jul 06, 2031				
	9770570	May 03, 2036		U-2148		
	9770570*PED	Nov 03, 2036				
<u>NITRIC OXIDE - INOMAX</u>						
N 020845 004	11931377	Jun 03, 2029	DS DP U-3903			
<u>NITRIC OXIDE - GENOSYL</u>						
N 202860 001	10124142	Aug 18, 2025		U-3037		
	10213572	Feb 12, 2036	DP			
	10737051	Oct 20, 2035	DP			
	10814092	Oct 17, 2025		U-3037		
	10926054	Aug 13, 2029	DP			
	11103669	Jun 21, 2030	DP			
	11291793	Aug 18, 2025	DP			
	11383059	Aug 18, 2025		U-3037		
	11511252	Sep 21, 2029	DP			
	11554241	Aug 18, 2025		U-3037		
	11672938	Jul 22, 2040		U-3037		
	7560076	Apr 21, 2027	DP			
	7618594	Oct 17, 2026	DP			
	7947227	Oct 17, 2026		U-3037		
	8057742	Jan 18, 2026		U-3037		
	8226916	Aug 18, 2025		U-3037		
	8607785	Jul 14, 2030	DP			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>NITRIC OXIDE - GENOSYL</u>						
N 202860	001 8609028	Aug 18, 2025	U-3037			
	8821801	Aug 18, 2025	DP			
	8944049	Aug 13, 2029	DP			
	9522249	Aug 18, 2025	DP			
	9604028	Aug 13, 2029	U-2793			
	9701538	Jan 28, 2029	DP			
	9956373	Aug 18, 2025	U-3037			
<u>NITROGLYCERIN - NITROLINGUAL PUMPSPRAY</u>						
N 018705	002 7872049	Mar 12, 2029	DP U-2223			
<u>NITROGLYCERIN - GONITRO</u>						
N 208424	001 9101592	Mar 11, 2032	DP			
<u>NOREPINEPHRINE BITARTRATE - NOREPINEPHRINE BITARTRATE IN 5% DEXTROSE</u>						
N 214313	001 12097170	Mar 08, 2041	DP U-3995			
<u>NOREPINEPHRINE BITARTRATE - NOREPINEPHRINE BITARTRATE IN 5% DEXTROSE</u>						
N 214313	002 12097170	Mar 08, 2041	DP U-3995			
<u>NOREPINEPHRINE BITARTRATE - NOREPINEPHRINE BITARTRATE IN 5% DEXTROSE</u>						
N 214313	003 12097170	Mar 08, 2041	DP U-3995			
<u>NOREPINEPHRINE BITARTRATE - NOREPINEPHRINE BITARTRATE IN 0.9% SODIUM CHLORIDE</u>						
N 214628	001 10159657	Jan 30, 2038	DP			
	10226436	Jan 30, 2038	DP U-3461			
	10420735	Jan 30, 2038	DP U-3461			
	10568850	Jan 30, 2038	DP			
	11413259	Jan 30, 2038	DP U-3461			
	11602508	Jan 30, 2038	DP			
<u>NOREPINEPHRINE BITARTRATE - NOREPINEPHRINE BITARTRATE IN 0.9% SODIUM CHLORIDE</u>						
N 214628	002 10159657	Jan 30, 2038	DP			
	10226436	Jan 30, 2038	DP U-3461			
	10420735	Jan 30, 2038	DP U-3461			
	10568850	Jan 30, 2038	DP			
	11413259	Jan 30, 2038	DP U-3461			
	11602508	Jan 30, 2038	DP			
<u>NOREPINEPHRINE BITARTRATE - NOREPINEPHRINE BITARTRATE IN 0.9% SODIUM CHLORIDE</u>						
N 214628	003 10159657	Jan 30, 2038	DP			
	10226436	Jan 30, 2038	DP U-3461			
	10420735	Jan 30, 2038	DP U-3461			
	10568850	Jan 30, 2038	DP			
	11413259	Jan 30, 2038	DP U-3461			
	11602508	Jan 30, 2038	DP			
<u>NOREPINEPHRINE BITARTRATE - NOREPINEPHRINE BITARTRATE IN 0.9% SODIUM CHLORIDE</u>						
N 215700	001 10888534	Apr 26, 2039	DP			
<u>NOREPINEPHRINE BITARTRATE - NOREPINEPHRINE BITARTRATE IN 0.9% SODIUM CHLORIDE</u>						
N 215700	002 10888534	Apr 26, 2039	DP			
<u>NOREPINEPHRINE BITARTRATE - NOREPINEPHRINE BITARTRATE IN 0.9% SODIUM CHLORIDE</u>						
N 215700	003 10888534	Apr 26, 2039	DP			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>NORGESTREL - OPILL</u>						
N 017031	001				RTO	Jul 13, 2026
<u>NUSINERSEN SODIUM - SPINRAZA</u>						
N 209531	001	10266822	Dec 05, 2025	U-1942		
		10266822	Dec 05, 2025	U-1943		
		10266822	Dec 05, 2025	U-1944		
		10436802	Sep 11, 2035	U-1941		
		10436802	Sep 11, 2035	U-1942		
		10436802	Sep 11, 2035	U-1943		
		10436802	Sep 11, 2035	U-1944		
		10436802	Sep 11, 2035	U-2093		
		10436802	Sep 11, 2035	U-2094		
		12013403	Mar 04, 2036	U-1941		
		12013403	Mar 04, 2036	U-1942		
		12013403	Mar 04, 2036	U-1943		
		12013403	Mar 04, 2036	U-1944		
		12013403	Mar 04, 2036	U-2093		
		12013403	Mar 04, 2036	U-2094		
		7838657	Jul 11, 2027	DS		
		8110560	Dec 05, 2025	U-1942		
		8110560	Dec 05, 2025	U-1943		
		8110560	Dec 05, 2025	U-1944		
		8361977	Dec 23, 2030	DS DP		
		8980853	Nov 24, 2030	U-1941		
		9717750	Jun 17, 2030	U-1942		
		9717750	Jun 17, 2030	U-1943		
		9717750	Jun 17, 2030	U-2093		
		9717750	Jun 17, 2030	U-2094		
		9926559	Jan 09, 2034	U-1943		
<u>OBETICHOLIC ACID - OCALIVA</u>						
N 207999	001	10047117	Sep 06, 2033	U-1854		
		10052337	Apr 26, 2036	DP		
		10174073	Jun 17, 2033	DS		
		10751349	Apr 26, 2036	DP		
		10758549	Apr 26, 2036	U-2945		
		9238673	Jun 17, 2033	DP		
		RE48286	Feb 21, 2027	DS DP		
<u>OBETICHOLIC ACID - OCALIVA</u>						
N 207999	002	10047117	Sep 06, 2033	U-1854		
		10052337	Apr 26, 2036	DP		
		10174073	Jun 17, 2033	DS		
		10751349	Apr 26, 2036	DP		
		10758549	Apr 26, 2036	U-2945		
		9238673	Jun 17, 2033	DP		
		RE48286	Feb 21, 2027	DS DP		
<u>OCTREOTIDE ACETATE - MYCAPSSA</u>						
N 208232	001	10238709	Feb 03, 2036	U-2857	ODE-474	Jun 26, 2027
		10695397	Feb 03, 2036	U-2857		
		11052126	Feb 03, 2036	U-2857		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>OCTREOTIDE ACETATE - MYCAPSSA</u>						
N 208232	001	11141457	Dec 28, 2040	U-3232		
		11338011	Feb 03, 2036	U-2857		
		11510963	Feb 03, 2036	U-2857		
		11857595	Feb 03, 2036	U-3784		
		11890316	Dec 28, 2040	U-3232		
		11969471	Sep 17, 2029	DP		
		11986529	Sep 17, 2029	U-2857		
		8329198	Sep 17, 2029	DP		
		8535695	Sep 17, 2029	U-2857		
		9265812	Sep 17, 2029	DP		
		9566246	Sep 17, 2029	DP		
<u>OCTREOTIDE ACETATE - BYNFEZIA PEN</u>						
N 213224	001	10342850	May 15, 2038	DP		
		11052196	Nov 03, 2040	U-4007		
		11052196	Nov 03, 2040	U-4008		
		11052196	Nov 03, 2040	U-4009		
		11246991	Nov 03, 2040	U-4007		
		11246991	Nov 03, 2040	U-4008		
		11246991	Nov 03, 2040	U-4009		
		11534553	Nov 03, 2040	U-4007		
		11534553	Nov 03, 2040	U-4008		
		11534553	Nov 03, 2040	U-4009		
<u>ODEVIXIBAT - BYLVAY</u>						
N 215498	001	10011633	Nov 08, 2031	U-3186	I-918	Jun 13, 2026
		10011633	Nov 08, 2031	U-3187	NCE	Jul 20, 2026
		10011633	Nov 08, 2031	U-3648	ODE-363	Jul 20, 2028
		10011633	Nov 08, 2031	U-3649	ODE-436	Jun 13, 2030
		10093697	Nov 08, 2031	U-3186		
		10093697	Nov 08, 2031	U-3187		
		10093697	Nov 08, 2031	U-3648		
		10093697	Nov 08, 2031	U-3649		
		10487111	Nov 08, 2031	U-3186		
		10487111	Nov 08, 2031	U-3187		
		10487111	Nov 08, 2031	U-3648		
		10487111	Nov 08, 2031	U-3649		
		10975046	Jun 20, 2039	DS		
		10981952	Nov 08, 2031	U-3186		
		10981952	Nov 08, 2031	U-3187		
		10981952	Nov 08, 2031	U-3648		
		10981952	Nov 08, 2031	U-3649		
		11365182	Jun 20, 2039	U-3186		
		11365182	Jun 20, 2039	U-3187		
		11365182	Jun 20, 2039	U-3648		
		11365182	Jun 20, 2039	U-3649		
		11583539	Nov 12, 2041	U-3186		
		11732006	Nov 08, 2031	U-3186		
		11732006	Nov 08, 2031	U-3187		
		11732006	Nov 08, 2031	U-3648		
		11732006	Nov 08, 2031	U-3649		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>ODEVIXIBAT - BYLVAY</u>						
N 215498	001	11801226	Jun 20, 2039	DP		
		11802115	Jun 20, 2039	DP		
		12091394	Jun 20, 2039	DS		
		9694018	Nov 08, 2031		U-3186	
		9694018	Nov 08, 2031		U-3187	
		9694018	Nov 08, 2031		U-3648	
		9694018	Nov 08, 2031		U-3649	
<u>ODEVIXIBAT - BYLVAY</u>						
N 215498	002	10011633	Nov 08, 2031		I-918	Jun 13, 2026
		10011633	Nov 08, 2031		NCE	Jul 20, 2026
		10011633	Nov 08, 2031		ODE-363	Jul 20, 2028
		10011633	Nov 08, 2031		ODE-436	Jun 13, 2030
		10093697	Nov 08, 2031			
		10093697	Nov 08, 2031		U-3186	
		10093697	Nov 08, 2031		U-3187	
		10093697	Nov 08, 2031		U-3648	
		10093697	Nov 08, 2031		U-3649	
		10487111	Nov 08, 2031		U-3186	
		10487111	Nov 08, 2031		U-3187	
		10487111	Nov 08, 2031		U-3648	
		10487111	Nov 08, 2031		U-3649	
		10975046	Jun 20, 2039	DS		
		10981952	Nov 08, 2031		U-3186	
		10981952	Nov 08, 2031		U-3187	
		10981952	Nov 08, 2031		U-3648	
		10981952	Nov 08, 2031		U-3649	
		11365182	Jun 20, 2039		U-3186	
		11365182	Jun 20, 2039		U-3187	
		11365182	Jun 20, 2039		U-3648	
		11365182	Jun 20, 2039		U-3649	
		11583539	Nov 12, 2041		U-3186	
		11732006	Nov 08, 2031		U-3186	
		11732006	Nov 08, 2031		U-3187	
		11732006	Nov 08, 2031		U-3648	
		11732006	Nov 08, 2031		U-3649	
		11801226	Jun 20, 2039	DP		
		11802115	Jun 20, 2039	DP		
		12091394	Jun 20, 2039	DS		
		9694018	Nov 08, 2031		U-3186	
		9694018	Nov 08, 2031		U-3187	
		9694018	Nov 08, 2031		U-3648	
		9694018	Nov 08, 2031		U-3649	
<u>ODEVIXIBAT - BYLVAY</u>						
N 215498	003	10011633	Nov 08, 2031		I-918	Jun 13, 2026
		10011633	Nov 08, 2031		NCE	Jul 20, 2026
		10011633	Nov 08, 2031		ODE-363	Jul 20, 2028
		10011633	Nov 08, 2031		ODE-436	Jun 13, 2030
		10093697	Nov 08, 2031			
		10093697	Nov 08, 2031		U-3186	
		10093697	Nov 08, 2031		U-3187	
		10093697	Nov 08, 2031		U-3648	

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>ODEVIXIBAT - BYLVAY</u>						
N 215498 003	10093697	Nov 08, 2031	U-3649			
	10487111	Nov 08, 2031	U-3186			
	10487111	Nov 08, 2031	U-3187			
	10487111	Nov 08, 2031	U-3648			
	10487111	Nov 08, 2031	U-3649			
	10975046	Jun 20, 2039	DS			
	10981952	Nov 08, 2031	U-3186			
	10981952	Nov 08, 2031	U-3187			
	10981952	Nov 08, 2031	U-3648			
	10981952	Nov 08, 2031	U-3649			
	11365182	Jun 20, 2039	U-3186			
	11365182	Jun 20, 2039	U-3187			
	11365182	Jun 20, 2039	U-3648			
	11365182	Jun 20, 2039	U-3649			
	11583539	Nov 12, 2041	U-3186			
	11732006	Nov 08, 2031	U-3186			
	11732006	Nov 08, 2031	U-3187			
	11732006	Nov 08, 2031	U-3648			
	11732006	Nov 08, 2031	U-3649			
	11801226	Jun 20, 2039	DP			
	11802115	Jun 20, 2039	DP			
	12091394	Jun 20, 2039	DS			
	9694018	Nov 08, 2031	U-3186			
	9694018	Nov 08, 2031	U-3187			
	9694018	Nov 08, 2031	U-3648			
	9694018	Nov 08, 2031	U-3649			
<u>ODEVIXIBAT - BYLVAY</u>						
N 215498 004	10011633	Nov 08, 2031	U-3186		I-918	Jun 13, 2026
	10011633	Nov 08, 2031	U-3187		NCE	Jul 20, 2026
	10011633	Nov 08, 2031	U-3648		ODE-363	Jul 20, 2028
	10011633	Nov 08, 2031	U-3649		ODE-436	Jun 13, 2030
	10093697	Nov 08, 2031	U-3186			
	10093697	Nov 08, 2031	U-3187			
	10093697	Nov 08, 2031	U-3648			
	10093697	Nov 08, 2031	U-3649			
	10487111	Nov 08, 2031	U-3186			
	10487111	Nov 08, 2031	U-3187			
	10487111	Nov 08, 2031	U-3648			
	10487111	Nov 08, 2031	U-3649			
	10975046	Jun 20, 2039	DS			
	10981952	Nov 08, 2031	U-3186			
	10981952	Nov 08, 2031	U-3187			
	10981952	Nov 08, 2031	U-3648			
	10981952	Nov 08, 2031	U-3649			
	11365182	Jun 20, 2039	U-3186			
	11365182	Jun 20, 2039	U-3187			
	11365182	Jun 20, 2039	U-3648			
	11365182	Jun 20, 2039	U-3649			
	11583539	Nov 12, 2041	U-3186			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>ODEVIXIBAT - BYLVAY</u>						
N 215498	004	11732006	Nov 08, 2031	U-3186		
		11732006	Nov 08, 2031	U-3187		
		11732006	Nov 08, 2031	U-3648		
		11732006	Nov 08, 2031	U-3649		
		11801226	Jun 20, 2039	DP		
		11802115	Jun 20, 2039	DP		
		12091394	Jun 20, 2039	DS		
		9694018	Nov 08, 2031	U-3186		
		9694018	Nov 08, 2031	U-3187		
		9694018	Nov 08, 2031	U-3648		
		9694018	Nov 08, 2031	U-3649		
<u>OLANZAPINE; SAMIDORPHAN L-MALATE - LYBALVI</u>						
N 213378	001	10300054	Aug 23, 2031	DP U-3140	NCE	May 28, 2026
		10300054	Aug 23, 2031	DP U-3141		
		10716785	Aug 23, 2031	U-3136		
		10716785	Aug 23, 2031	U-3137		
		11185541	Aug 23, 2031	U-3140		
		11241425	Aug 23, 2031	U-3137		
		11351166	Aug 23, 2031	U-3140		
		11351166	Aug 23, 2031	U-3141		
		11707466	Nov 12, 2041	DP		
		11793805	Aug 23, 2031	U-3734		
		11951111	Nov 12, 2041	U-3886		
		11951111	Nov 12, 2041	U-3887		
		7262298	Nov 23, 2025	DS		
		8778960	Feb 13, 2032	U-3136		
		8778960	Feb 13, 2032	U-3137		
		9119848	Aug 30, 2031	DS		
		9126977	Aug 23, 2031	DP U-3136		
		9126977	Aug 23, 2031	DP U-3137		
		9517235	Aug 23, 2031	U-3138		
		9517235	Aug 23, 2031	U-3139		
<u>OLANZAPINE; SAMIDORPHAN L-MALATE - LYBALVI</u>						
N 213378	002	10300054	Aug 23, 2031	DP U-3140	NCE	May 28, 2026
		10300054	Aug 23, 2031	DP U-3141		
		10716785	Aug 23, 2031	U-3136		
		10716785	Aug 23, 2031	U-3137		
		11185541	Aug 23, 2031	U-3140		
		11241425	Aug 23, 2031	U-3137		
		11351166	Aug 23, 2031	U-3140		
		11351166	Aug 23, 2031	U-3141		
		11707466	Nov 12, 2041	DP		
		11793805	Aug 23, 2031	U-3734		
		11951111	Nov 12, 2041	U-3886		
		11951111	Nov 12, 2041	U-3887		
		7262298	Nov 23, 2025	DS		
		8778960	Feb 13, 2032	U-3136		
		8778960	Feb 13, 2032	U-3137		
		9119848	Aug 30, 2031	DS		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>OLANZAPINE; SAMIDORPHAN L-MALATE - LYBALVI</u>						
N 213378 002	9126977	Aug 23, 2031	DP U-3136			
	9126977	Aug 23, 2031	DP U-3137			
	9517235	Aug 23, 2031	U-3138			
	9517235	Aug 23, 2031	U-3139			
<u>OLANZAPINE; SAMIDORPHAN L-MALATE - LYBALVI</u>						
N 213378 003	10300054	Aug 23, 2031	DP U-3140		NCE	May 28, 2026
	10300054	Aug 23, 2031	DP U-3141			
	10716785	Aug 23, 2031	U-3136			
	10716785	Aug 23, 2031	U-3137			
	11185541	Aug 23, 2031	U-3140			
	11241425	Aug 23, 2031	U-3137			
	11351166	Aug 23, 2031	U-3140			
	11351166	Aug 23, 2031	U-3141			
	11707466	Nov 12, 2041	DP			
	11793805	Aug 23, 2031	U-3734			
	11951111	Nov 12, 2041	U-3886			
	11951111	Nov 12, 2041	U-3887			
	7262298	Nov 23, 2025	DS			
	8778960	Feb 13, 2032	U-3136			
	8778960	Feb 13, 2032	U-3137			
	9119848	Aug 30, 2031	DS			
	9126977	Aug 23, 2031	DP U-3136			
	9126977	Aug 23, 2031	DP U-3137			
	9517235	Aug 23, 2031	U-3138			
	9517235	Aug 23, 2031	U-3139			
<u>OLANZAPINE; SAMIDORPHAN L-MALATE - LYBALVI</u>						
N 213378 004	10300054	Aug 23, 2031	DP U-3140		NCE	May 28, 2026
	10300054	Aug 23, 2031	DP U-3141			
	10716785	Aug 23, 2031	U-3136			
	10716785	Aug 23, 2031	U-3137			
	11185541	Aug 23, 2031	U-3140			
	11241425	Aug 23, 2031	U-3137			
	11351166	Aug 23, 2031	U-3140			
	11351166	Aug 23, 2031	U-3141			
	11707466	Nov 12, 2041	DP			
	11793805	Aug 23, 2031	U-3734			
	11951111	Nov 12, 2041	U-3886			
	11951111	Nov 12, 2041	U-3887			
	7262298	Nov 23, 2025	DS			
	8778960	Feb 13, 2032	U-3136			
	8778960	Feb 13, 2032	U-3137			
	9119848	Aug 30, 2031	DS			
	9126977	Aug 23, 2031	DP U-3136			
	9126977	Aug 23, 2031	DP U-3137			
	9517235	Aug 23, 2031	U-3138			
	9517235	Aug 23, 2031	U-3139			
<u>OLAPARIB - LYNPARZA</u>						
N 206162 001	8143241	Aug 12, 2027	U-1634			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>OLAPARIB - LYNPARZA</u>						
N 206162 001	8247416	Sep 24, 2028	DS			
	8859562	Aug 04, 2031		U-1634		
<u>OLAPARIB - LYNPARZA</u>						
N 208558 001	11633396	Oct 07, 2029	DP		I-885	Mar 11, 2025
	11970530	Oct 25, 2041		U-3929	I-914	May 31, 2026
	11970530	Oct 25, 2041		U-3930	ODE-226	Dec 19, 2025
	11970530	Oct 25, 2041		U-3931	ODE-283	Dec 27, 2026
	11975001	Oct 07, 2029	DP		ODE-306	May 08, 2027
	12048695	Oct 07, 2029	DP			
	12144810	Oct 07, 2029	DP			
	7449464	Sep 08, 2027	DS DP			
	8071579	Aug 12, 2027		U-2480		
	8071579	Aug 12, 2027		U-2482		
	8071579	Aug 12, 2027		U-2483		
	8071579	Aug 12, 2027		U-2716		
	8071579	Aug 12, 2027		U-2819		
	8071579	Aug 12, 2027		U-2820		
	8071579	Aug 12, 2027		U-2821		
	8071579	Aug 12, 2027		U-2822		
	8071579	Aug 12, 2027		U-2823		
	8071579	Aug 12, 2027		U-2824		
	8071579	Aug 12, 2027		U-2832		
	8071579	Aug 12, 2027		U-2833		
	8071579	Aug 12, 2027		U-3333		
	8071579	Aug 12, 2027		U-3631		
	8071579	Aug 12, 2027		U-3695		
	8143241	Aug 12, 2027		U-2480		
	8143241	Aug 12, 2027		U-2482		
	8143241	Aug 12, 2027		U-2483		
	8143241	Aug 12, 2027		U-2716		
	8143241	Aug 12, 2027		U-2819		
	8143241	Aug 12, 2027		U-2820		
	8143241	Aug 12, 2027		U-2821		
	8143241	Aug 12, 2027		U-2822		
	8143241	Aug 12, 2027		U-2823		
	8143241	Aug 12, 2027		U-2824		
	8143241	Aug 12, 2027		U-2832		
	8143241	Aug 12, 2027		U-2833		
	8143241	Aug 12, 2027		U-3333		
	8143241	Aug 12, 2027		U-3631		
	8143241	Aug 12, 2027		U-3695		
	8475842	Dec 31, 2029	DP			
	8859562	Aug 04, 2031		U-2101		
	8859562	Aug 04, 2031		U-2103		
	8859562	Aug 04, 2031		U-2480		
	8859562	Aug 04, 2031		U-2482		
	8859562	Aug 04, 2031		U-2483		
	8859562	Aug 04, 2031		U-2716		
	8859562	Aug 04, 2031		U-2819		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>OLAPARIB - LYNPARZA</u>						
N 208558 001	8859562	Aug 04, 2031	U-2820			
	8859562	Aug 04, 2031	U-2821			
	8859562	Aug 04, 2031	U-2822			
	8859562	Aug 04, 2031	U-2823			
	8859562	Aug 04, 2031	U-2824			
	8859562	Aug 04, 2031	U-2832			
	8859562	Aug 04, 2031	U-2833			
	8859562	Aug 04, 2031	U-3333			
	8859562	Aug 04, 2031	U-3631			
	8859562	Aug 04, 2031	U-3695			
<u>OLAPARIB - LYNPARZA</u>						
N 208558 002	11633396	Oct 07, 2029	DP		I-885	Mar 11, 2025
	11970530	Oct 25, 2041	U-3929		I-914	May 31, 2026
	11970530	Oct 25, 2041	U-3930		ODE-226	Dec 19, 2025
	11970530	Oct 25, 2041	U-3931		ODE-283	Dec 27, 2026
	11975001	Oct 07, 2029	DP		ODE-306	May 08, 2027
	12048695	Oct 07, 2029	DP			
	12144810	Oct 07, 2029	DP			
	7449464	Sep 08, 2027	DS DP			
	8071579	Aug 12, 2027	U-2480			
	8071579	Aug 12, 2027	U-2482			
	8071579	Aug 12, 2027	U-2483			
	8071579	Aug 12, 2027	U-2716			
	8071579	Aug 12, 2027	U-2819			
	8071579	Aug 12, 2027	U-2820			
	8071579	Aug 12, 2027	U-2821			
	8071579	Aug 12, 2027	U-2822			
	8071579	Aug 12, 2027	U-2823			
	8071579	Aug 12, 2027	U-2824			
	8071579	Aug 12, 2027	U-2832			
	8071579	Aug 12, 2027	U-2833			
	8071579	Aug 12, 2027	U-3333			
	8071579	Aug 12, 2027	U-3631			
	8071579	Aug 12, 2027	U-3695			
	8143241	Aug 12, 2027	U-2480			
	8143241	Aug 12, 2027	U-2482			
	8143241	Aug 12, 2027	U-2483			
	8143241	Aug 12, 2027	U-2716			
	8143241	Aug 12, 2027	U-2819			
	8143241	Aug 12, 2027	U-2820			
	8143241	Aug 12, 2027	U-2821			
	8143241	Aug 12, 2027	U-2822			
	8143241	Aug 12, 2027	U-2823			
	8143241	Aug 12, 2027	U-2824			
	8143241	Aug 12, 2027	U-2832			
	8143241	Aug 12, 2027	U-2833			
	8143241	Aug 12, 2027	U-3333			
	8143241	Aug 12, 2027	U-3631			
	8143241	Aug 12, 2027	U-3695			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>OLAPARIB - LYNPARZA</u>						
N 208558	002	8475842	Dec 31, 2029	DP		
		8859562	Aug 04, 2031	U-2101		
		8859562	Aug 04, 2031	U-2103		
		8859562	Aug 04, 2031	U-2480		
		8859562	Aug 04, 2031	U-2482		
		8859562	Aug 04, 2031	U-2483		
		8859562	Aug 04, 2031	U-2716		
		8859562	Aug 04, 2031	U-2819		
		8859562	Aug 04, 2031	U-2820		
		8859562	Aug 04, 2031	U-2821		
		8859562	Aug 04, 2031	U-2822		
		8859562	Aug 04, 2031	U-2823		
		8859562	Aug 04, 2031	U-2824		
		8859562	Aug 04, 2031	U-2832		
		8859562	Aug 04, 2031	U-2833		
		8859562	Aug 04, 2031	U-3333		
		8859562	Aug 04, 2031	U-3631		
		8859562	Aug 04, 2031	U-3695		
<u>OLICERIDINE - OLINVYK</u>						
N 210730	001	11077098	Mar 23, 2032	DS DP U-2986	NCE	Oct 30, 2025
		11931350	Mar 23, 2032	DS U-2986		
		8835488	Mar 23, 2032	DS DP U-2986		
		9309234	Mar 23, 2032	DS DP U-2986		
		9642842	Mar 23, 2032	DP U-2986		
<u>OLICERIDINE - OLINVYK</u>						
N 210730	002	11077098	Mar 23, 2032	DS DP U-2986	NCE	Oct 30, 2025
		11931350	Mar 23, 2032	DS U-2986		
		8835488	Mar 23, 2032	DS DP U-2986		
		9309234	Mar 23, 2032	DS DP U-2986		
		9642842	Mar 23, 2032	DP U-2986		
<u>OLICERIDINE - OLINVYK</u>						
N 210730	003	11077098	Mar 23, 2032	DS DP U-2986	NCE	Oct 30, 2025
		11931350	Mar 23, 2032	DS U-2986		
		8835488	Mar 23, 2032	DS DP U-2986		
		9309234	Mar 23, 2032	DS DP U-2986		
		9642842	Mar 23, 2032	DP U-2986		
<u>OLIVE OIL; SOYBEAN OIL - CLINOLIPID 20%</u>						
N 204508	001				NPP	Apr 24, 2027
<u>OLODATEROL HYDROCHLORIDE - STRIVERDI RESPIMAT</u>						
N 203108	001	7220742	May 12, 2025	DS DP U-1547		
		7396341	Oct 10, 2026	DP U-1547		
		7727984	Jan 19, 2027	DS		
		7837235	Mar 13, 2028	DP		
		7896264	May 26, 2025	DP		
		8034809	May 12, 2025	U-1547		
		8733341	Oct 16, 2030	DP		
		9027967	Mar 31, 2027	DP		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>OLODATEROL HYDROCHLORIDE; TIOTROPIUM BROMIDE - STIOLTO RESPIMAT</u>						
N 206756	001 7220742	May 12, 2025	DS DP U-1703			
	7284474*PED	Feb 26, 2025				
	7396341	Oct 10, 2026	DP			
	7396341*PED	Apr 10, 2027				
	7727984	Jan 19, 2027	DS			
	7837235	Mar 13, 2028	DP			
	7837235*PED	Sep 13, 2028				
	7896264	May 26, 2025	DP			
	8034809	May 12, 2025	U-1702			
	8733341	Oct 16, 2030	DP			
	9027967	Mar 31, 2027	DP			
<u>OLOPATADINE HYDROCHLORIDE - PATADAY ONCE DAILY RELIEF</u>						
N 206276	001 8791154	May 19, 2032	DP U-1680			
	9533053	May 19, 2032	DP			
<u>OLUTASIDENIB - REZLIDHIA</u>						
N 215814	001 10414752	Sep 18, 2035	DP		NCE	Dec 01, 2027
	10532047	May 16, 2039	DS		ODE-413	Dec 01, 2029
	10550098	Sep 18, 2035	DP			
	10959994	May 16, 2039	DP			
	11013733	May 16, 2039	U-3496			
	11013734	May 16, 2039	U-3495			
	11376246	May 16, 2039	U-3495			
	11497743	May 16, 2039	U-3495			
	11498913	Sep 18, 2035	DP			
	11723905	Nov 12, 2039	DP			
	11738018	Jul 17, 2039	U-3684			
	12053463	May 16, 2039	DS DP			
	9834539	Sep 18, 2035	DS DP U-3497			
<u>OMACETAXINE MEPESUCCINATE - SYNRIBO</u>						
N 203585	001 6987103	Oct 26, 2026	U-1300			
<u>OMADACYCLINE TOSYLATE - NUZYRA</u>						
N 209816	001 10111890	Aug 03, 2037	U-2444		NCE	Oct 02, 2023
	10124014	Mar 05, 2029	U-2449		GAIN	Oct 02, 2028
	10383884	Oct 31, 2037	U-2576			
	10835542	Oct 31, 2037	U-2576			
	7553828	Jun 02, 2025	DS			
	8383610	Sep 23, 2030	DS			
	9265740	Mar 05, 2029	U-1569			
	9314475	Mar 18, 2031	DP			
	9724358	Mar 05, 2029	U-1569			
<u>OMADACYCLINE TOSYLATE - NUZYRA</u>						
N 209817	001 10124014	Mar 05, 2029	U-2449		NCE	Oct 02, 2023
	10383884	Oct 31, 2037	U-2576		GAIN	Oct 02, 2028
	10835542	Oct 31, 2037	U-2576			
	7553828	Jun 02, 2025	DS			
	9265740	Mar 05, 2029	DP			
	9724358	Mar 05, 2029	U-1569			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>OMAVELOXOLONE - SKYCLARYS</u>						
N 216718	001 11091430	Apr 20, 2029	U-3552		NCE	Feb 28, 2028
	11919838	Apr 20, 2029	U-3862		ODE-427	Feb 28, 2030
	11919838	Apr 20, 2029	U-3863			
	8124799	Dec 03, 2029	DS			
	8440854	Apr 20, 2029	DP			
	8993640	Apr 24, 2033	DS DP			
	9670147	Apr 20, 2029	DS			
	9701709	Apr 24, 2033	DS DP			
<u>OMBITASVIR; PARITAPREVR; RITONAVIR - TECHNIVIE</u>						
N 207931	001 8268349*PED	Feb 25, 2025				
	8399015*PED	Feb 25, 2025				
	8420596	Apr 10, 2031	DS DP			
	8420596*PED	Oct 10, 2031				
	8642538	Sep 10, 2029	DS DP	U-1638		
	8686026	Jun 09, 2031	DP			
	8691938	Apr 13, 2032	DS DP			
	9006387	Jun 10, 2030		U-1687		
	9044480	Apr 10, 2031		U-1638		
<u>OMEGA-3-CARBOXYLIC ACIDS - EPANOVA</u>						
N 205060	001 10117844	Jan 04, 2033		U-2447		
	7960370	Dec 20, 2026	DP			
	8383678	Feb 07, 2025	DP	U-1511		
	9012501	Feb 07, 2025	DP	U-1511		
	9050308	Jan 04, 2033		U-1511		
	9050309	Jan 04, 2033	DS			
	9132112	Feb 07, 2025	DP	U-1511		
<u>OMEPRAZOLE - OMEPRAZOLE</u>						
N 022032	001 9023391	Aug 16, 2025	DP			
<u>OMEPRAZOLE - OMEPRAZOLE</u>						
N 209400	001 10076494	Dec 08, 2036	DP			
	10835488	Dec 08, 2036	DP			
<u>OMEPRAZOLE; SODIUM BICARBONATE - KONVOMEF</u>						
N 213593	001 10751333	Jul 16, 2039	DP			
	11103492	Jul 16, 2039	DP			
	11633478	Jul 16, 2039	DP			
	11771686	Mar 01, 2040		U-623		
	11911473	Jul 16, 2039		U-623		
	11911473	Jul 16, 2039		U-624		
	12042539	Jul 16, 2039	DP			
<u>OMIDENEPAG ISOPROPYL - OMLONTI</u>						
N 215092	001 10179127	Jan 08, 2035	DP	U-3454	NCE	Sep 22, 2027
	10702511	Jan 08, 2035	DP	U-3454		
	10765750	Jan 08, 2035	DP			
	10774072	Jun 10, 2035	DS			
	11197849	Jan 08, 2035	DP	U-3454		
	11666563	Jul 16, 2039		U-3454		
	11793798	Jan 08, 2035	DP	U-3454		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>OMIDENEPAG ISOPROPYL - OMLONTI</u>						
N 215092	001	8648097	Oct 13, 2029	DS DP		
		8685986	Oct 13, 2029	DP		
		9415038	Jan 08, 2035	DP U-3454		
		RE48183	Jan 08, 2035	DP U-3454		
<u>ONDANSETRON - ZUPLENZ</u>						
N 022524	001	8580830	Nov 23, 2029	DP		
		9095577	Jul 13, 2030	DP		
<u>ONDANSETRON - ZUPLENZ</u>						
N 022524	002	8580830	Nov 23, 2029	DP		
		9095577	Jul 13, 2030	DP		
<u>OPICAPONE - ONGENTYS</u>						
N 212489	001	10071085	Mar 31, 2030	DP	NCE	Apr 24, 2025
		10583130	Mar 31, 2030	U-3869		
		12129247	Jan 11, 2032	U-4038		
		8168793	Apr 02, 2029	DS DP U-3869		
		8524746	Jul 14, 2029	U-3869		
		8907099	May 12, 2027	DS		
		9550759	Jul 26, 2026	U-3870		
		9550759	Jul 26, 2026	U-3871		
		9550759	Jul 26, 2026	U-3872		
		9630955	Dec 12, 2032	DS DP U-3869		
		9745290	Oct 10, 2027	DP U-3869		
<u>OPICAPONE - ONGENTYS</u>						
N 212489	002	10071085	Mar 31, 2030	DP	NCE	Apr 24, 2025
		10583130	Mar 31, 2030	U-3869		
		12129247	Jan 11, 2032	U-4038		
		8168793	Apr 02, 2029	DS DP U-3869		
		8524746	Jul 14, 2029	U-3869		
		8907099	May 12, 2027	DS		
		9550759	Jul 26, 2026	U-3870		
		9550759	Jul 26, 2026	U-3871		
		9550759	Jul 26, 2026	U-3872		
		9630955	Dec 12, 2032	DS DP U-3869		
		9745290	Oct 10, 2027	DP U-3869		
<u>ORITAVANCIN DIPHOSPHATE - ORBACTIV</u>						
N 206334	001	8420592	Aug 29, 2029	U-1570		
		9649352	Jul 16, 2035	DP		
		9682061	Apr 26, 2030	U-1569		
<u>ORITAVANCIN DIPHOSPHATE - KIMYRSA</u>						
N 214155	001	8420592	Aug 29, 2029	U-3101		
		9649352	Jul 16, 2035	DS DP		
		9682061	Apr 26, 2030	U-3101		
<u>OSILODROSTAT PHOSPHATE - ISTURISA</u>						
N 212801	001	10143680	Jul 06, 2035	DP	NCE	Mar 06, 2025
		10709691	Oct 12, 2035	U-2770	ODE-286	Mar 06, 2027
		8314097	Mar 27, 2029	DS DP		
		8609862	Jan 13, 2031	U-2770		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>OSILODROSTAT PHOSPHATE - ISTURISA</u>						
N 212801 001	8835646	Aug 23, 2026	DS DP			
	9434754	Jan 13, 2031	DS			
<u>OSILODROSTAT PHOSPHATE - ISTURISA</u>						
N 212801 002	10143680	Jul 06, 2035	DP		NCE	Mar 06, 2025
	10709691	Oct 12, 2035	U-2770		ODE-286	Mar 06, 2027
	8314097	Mar 27, 2029	DS DP			
	8609862	Jan 13, 2031	U-2770			
	8835646	Aug 23, 2026	DS DP			
	9434754	Jan 13, 2031	DS			
<u>OSILODROSTAT PHOSPHATE - ISTURISA</u>						
N 212801 003	10143680	Jul 06, 2035	DP		NCE	Mar 06, 2025
	10709691	Oct 12, 2035	U-2770		ODE-286	Mar 06, 2027
	8314097	Mar 27, 2029	DS DP			
	8609862	Jan 13, 2031	U-2770			
	8835646	Aug 23, 2026	DS DP			
	9434754	Jan 13, 2031	DS			
<u>OSIMERTINIB MESYLATE - TAGRISSO</u>						
N 208065 001	10183020	Jan 02, 2035	DP U-1777		I-941	Feb 16, 2027
	10183020	Jan 02, 2035	DP U-2289		I-952	Sep 25, 2027
	10183020	Jan 02, 2035	DP U-3016		ODE-176	Apr 18, 2025
	10183020	Jan 02, 2035	DP U-3823		ODE-337	Dec 18, 2027
	10183020	Jan 02, 2035	DP U-4010			
	11524951	Jul 25, 2032	DS DP			
	8946235	Aug 08, 2032	DS DP U-1777			
	8946235	Aug 08, 2032	DS DP U-2289			
	8946235	Aug 08, 2032	DS DP U-3016			
	8946235	Aug 08, 2032	DS DP U-3823			
	8946235	Aug 08, 2032	DS DP U-4010			
	9732058	Jul 25, 2032	DS DP U-1777			
	9732058	Jul 25, 2032	DS DP U-2289			
	9732058	Jul 25, 2032	DS DP U-3016			
	9732058	Jul 25, 2032	DS DP U-3823			
	9732058	Jul 25, 2032	DS DP U-4010			
<u>OSIMERTINIB MESYLATE - TAGRISSO</u>						
N 208065 002	10183020	Jan 02, 2035	DP U-1777		I-941	Feb 16, 2027
	10183020	Jan 02, 2035	DP U-2289		I-952	Sep 25, 2027
	10183020	Jan 02, 2035	DP U-3016		ODE-176	Apr 18, 2025
	10183020	Jan 02, 2035	DP U-3823		ODE-337	Dec 18, 2027
	10183020	Jan 02, 2035	DP U-4010			
	11524951	Jul 25, 2032	DS DP			
	8946235	Aug 08, 2032	DS DP U-1777			
	8946235	Aug 08, 2032	DS DP U-2289			
	8946235	Aug 08, 2032	DS DP U-3016			
	8946235	Aug 08, 2032	DS DP U-3823			
	8946235	Aug 08, 2032	DS DP U-4010			
	9732058	Jul 25, 2032	DS DP U-1777			
	9732058	Jul 25, 2032	DS DP U-2289			
	9732058	Jul 25, 2032	DS DP U-3016			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>OSIMERTINIB MESYLATE - TAGRISSO</u>						
N 208065	002 9732058	Jul 25, 2032	DS DP U-3823			
	9732058	Jul 25, 2032	DS DP U-4010			
<u>OSPEMIFENE - OSPHENA</u>						
N 203505	001 6245819	Jul 21, 2025		U-1370		
	6245819	Jul 21, 2025		U-905		
	8236861	Aug 11, 2026		U-1369		
	8236861	Aug 11, 2026		U-1370		
	8236861	Aug 11, 2026		U-905		
	8642079	Jul 09, 2028	DP			
<u>OTESECONAZOLE - VIVJOA</u>						
N 215888	001 10414751	Mar 17, 2036	DS DP		NCE	Apr 26, 2027
	11247981	May 09, 2033		U-3366	GAIN	Apr 26, 2032
	8236962	Apr 22, 2031	DS DP			
	8754227	Apr 22, 2031		U-3366		
	9840492	Mar 17, 2036	DS DP			
<u>OXCARBAZEPINE - OXTELLAR XR</u>						
N 202810	001 10220042	Apr 13, 2027		U-2501		
	11166960	Apr 13, 2027	DP			
	11896599	Apr 13, 2027	DP			
	7722898	Apr 13, 2027	DP			
	7910131	Apr 13, 2027		U-2041		
	8617600	Apr 13, 2027	DP			
	8821930	Apr 13, 2027	DP			
	9119791	Apr 13, 2027		U-2041		
	9351975	Apr 13, 2027	DP			
	9370525	Apr 13, 2027	DP			
	9855278	Apr 13, 2027	DP			
<u>OXCARBAZEPINE - OXTELLAR XR</u>						
N 202810	002 10220042	Apr 13, 2027		U-2501		
	11166960	Apr 13, 2027	DP			
	11896599	Apr 13, 2027	DP			
	7722898	Apr 13, 2027	DP			
	7910131	Apr 13, 2027		U-2041		
	8617600	Apr 13, 2027	DP			
	8821930	Apr 13, 2027	DP			
	9119791	Apr 13, 2027		U-2041		
	9351975	Apr 13, 2027	DP			
	9370525	Apr 13, 2027	DP			
	9855278	Apr 13, 2027	DP			
<u>OXCARBAZEPINE - OXTELLAR XR</u>						
N 202810	003 10220042	Apr 13, 2027		U-2501		
	11166960	Apr 13, 2027	DP			
	11896599	Apr 13, 2027	DP			
	7722898	Apr 13, 2027	DP			
	7910131	Apr 13, 2027		U-2041		
	8617600	Apr 13, 2027	DP			
	8821930	Apr 13, 2027	DP			
	9119791	Apr 13, 2027		U-2041		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>OXCARBAZEPINE - OXTELLAR XR</u>						
N 202810	003 9351975	Apr 13, 2027	DP			
	9370525	Apr 13, 2027	DP			
	9855278	Apr 13, 2027	DP			
<u>OXYBUTYNYN CHLORIDE - GELNIQUE</u>						
N 022204	001 10449173	Nov 06, 2029	DP U-2637			
	8920392	Mar 26, 2031	U-1644			
	9259388	Nov 06, 2029	U-1644			
<u>OXYCODONE - XTAMPZA ER</u>						
N 208090	001 10004729	Dec 10, 2030	DP U-1556			
	10188644	Sep 02, 2036	DP U-1556			
	10646485	Sep 02, 2036	DP U-1556			
	10668060	Dec 10, 2030	DP U-1556			
	7399488	Mar 24, 2025	DP			
	7771707	Mar 24, 2025	DP			
	8449909	Mar 24, 2025	DP			
	8557291	Mar 21, 2025	DP			
	8758813	Jun 10, 2025	U-1556			
	9682075	Dec 10, 2030	DP U-1556			
	9737530	Sep 02, 2036	DP U-1556			
	9968598	Sep 02, 2036	DP U-1556			
<u>OXYCODONE - XTAMPZA ER</u>						
N 208090	002 10004729	Dec 10, 2030	DP U-1556			
	10188644	Sep 02, 2036	DP U-1556			
	10646485	Sep 02, 2036	DP U-1556			
	10668060	Dec 10, 2030	DP U-1556			
	7399488	Mar 24, 2025	DP			
	7771707	Mar 24, 2025	DP			
	8449909	Mar 24, 2025	DP			
	8557291	Mar 21, 2025	DP			
	8758813	Jun 10, 2025	U-1556			
	9682075	Dec 10, 2030	DP U-1556			
	9737530	Sep 02, 2036	DP U-1556			
	9968598	Sep 02, 2036	DP U-1556			
<u>OXYCODONE - XTAMPZA ER</u>						
N 208090	003 10004729	Dec 10, 2030	DP U-1556			
	10188644	Sep 02, 2036	DP U-1556			
	10646485	Sep 02, 2036	DP U-1556			
	10668060	Dec 10, 2030	DP U-1556			
	7399488	Mar 24, 2025	DP			
	7771707	Mar 24, 2025	DP			
	8449909	Mar 24, 2025	DP			
	8557291	Mar 21, 2025	DP			
	8758813	Jun 10, 2025	U-1556			
	9682075	Dec 10, 2030	DP U-1556			
	9737530	Sep 02, 2036	DP U-1556			
	9968598	Sep 02, 2036	DP U-1556			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>OXYCODONE - XTAMPZA ER</u>						
N 208090 004	10004729	Dec 10, 2030	DP U-1556			
	10188644	Sep 02, 2036	DP U-1556			
	10646485	Sep 02, 2036	DP U-1556			
	10668060	Dec 10, 2030	DP U-1556			
	7399488	Mar 24, 2025	DP			
	7771707	Mar 24, 2025	DP			
	8449909	Mar 24, 2025	DP			
	8557291	Mar 21, 2025	DP			
	8758813	Jun 10, 2025	U-1556			
	9682075	Dec 10, 2030	DP U-1556			
	9737530	Sep 02, 2036	DP U-1556			
	9968598	Sep 02, 2036	DP U-1556			
<u>OXYCODONE - XTAMPZA ER</u>						
N 208090 005	10004729	Dec 10, 2030	DP U-1556			
	10188644	Sep 02, 2036	DP U-1556			
	10646485	Sep 02, 2036	DP U-1556			
	10668060	Dec 10, 2030	DP U-1556			
	7399488	Mar 24, 2025	DP			
	7771707	Mar 24, 2025	DP			
	8449909	Mar 24, 2025	DP			
	8557291	Mar 21, 2025	DP			
	8758813	Jun 10, 2025	U-1556			
	9682075	Dec 10, 2030	DP U-1556			
	9737530	Sep 02, 2036	DP U-1556			
	9968598	Sep 02, 2036	DP U-1556			
<u>OXYCODONE HYDROCHLORIDE - OXYCONTIN</u>						
N 022272 001	10407434	Mar 30, 2025	DS			
	10696684	Mar 30, 2025	DS			
	11304908	Aug 24, 2027	DP			
	11304909	Aug 24, 2027	U-1556			
	11964056	Aug 24, 2027	DP			
	12060361	Mar 30, 2025	DP U-1556			
	8808741	Aug 24, 2027	U-1556			
	8894987	Mar 29, 2030	DP			
	8894988	Aug 24, 2027	DP			
	9073933	Mar 30, 2025	DS			
	9492389	Aug 24, 2027	DP			
	9492391	Aug 24, 2027	U-1556			
	9492392	Aug 24, 2027	DP			
	9492393	Aug 24, 2027	U-1556			
	9522919	Mar 30, 2025	DS DP			
	9763933	Aug 24, 2027	DP			
	9770416	Aug 24, 2027	DP			
	9775808	Aug 24, 2027	DP			
<u>OXYCODONE HYDROCHLORIDE - OXYCONTIN</u>						
N 022272 002	10407434	Mar 30, 2025	DS			
	10696684	Mar 30, 2025	DS			
	11304908	Aug 24, 2027	DP			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>OXYCODONE HYDROCHLORIDE - OXYCONTIN</u>						
N 022272 002	11304909	Aug 24, 2027	U-1556			
	11964056	Aug 24, 2027	DP			
	12060361	Mar 30, 2025	DP U-1556			
	8808741	Aug 24, 2027	U-1556			
	8894987	Mar 29, 2030	DP			
	8894988	Aug 24, 2027	DP			
	9073933	Mar 30, 2025	DS			
	9492389	Aug 24, 2027	DP			
	9492391	Aug 24, 2027	U-1556			
	9492392	Aug 24, 2027	DP			
	9492393	Aug 24, 2027	U-1556			
	9522919	Mar 30, 2025	DS DP			
	9763933	Aug 24, 2027	DP			
	9770416	Aug 24, 2027	DP			
	9775808	Aug 24, 2027	DP			
<u>OXYCODONE HYDROCHLORIDE - OXYCONTIN</u>						
N 022272 003	10407434	Mar 30, 2025	DS			
	10696684	Mar 30, 2025	DS			
	11304908	Aug 24, 2027	DP			
	11304909	Aug 24, 2027	U-1556			
	11964056	Aug 24, 2027	DP			
	12060361	Mar 30, 2025	DP U-1556			
	8808741	Aug 24, 2027	U-1556			
	8894987	Mar 29, 2030	DP			
	8894988	Aug 24, 2027	DP			
	9073933	Mar 30, 2025	DS			
	9492389	Aug 24, 2027	DP			
	9492391	Aug 24, 2027	U-1556			
	9492392	Aug 24, 2027	DP			
	9492393	Aug 24, 2027	U-1556			
	9522919	Mar 30, 2025	DS DP			
	9763933	Aug 24, 2027	DP			
	9770416	Aug 24, 2027	DP			
	9775808	Aug 24, 2027	DP			
<u>OXYCODONE HYDROCHLORIDE - OXYCONTIN</u>						
N 022272 004	10407434	Mar 30, 2025	DS			
	10696684	Mar 30, 2025	DS			
	11304908	Aug 24, 2027	DP			
	11304909	Aug 24, 2027	U-1556			
	11964056	Aug 24, 2027	DP			
	12060361	Mar 30, 2025	DP U-1556			
	8808741	Aug 24, 2027	U-1556			
	8894987	Mar 29, 2030	DP			
	8894988	Aug 24, 2027	DP			
	9073933	Mar 30, 2025	DS			
	9492389	Aug 24, 2027	DP			
	9492391	Aug 24, 2027	U-1556			
	9492392	Aug 24, 2027	DP			
	9492393	Aug 24, 2027	U-1556			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>OXYCODONE HYDROCHLORIDE - OXYCONTIN</u>						
N 022272 004	9522919	Mar 30, 2025	DS DP			
	9763933	Aug 24, 2027	DP			
	9770416	Aug 24, 2027	DP			
	9775808	Aug 24, 2027	DP			
<u>OXYCODONE HYDROCHLORIDE - OXYCONTIN</u>						
N 022272 005	10407434	Mar 30, 2025	DS			
	10696684	Mar 30, 2025	DS			
	11304908	Aug 24, 2027	DP			
	11304909	Aug 24, 2027		U-1556		
	11964056	Aug 24, 2027	DP			
	12060361	Mar 30, 2025	DP	U-1556		
	8808741	Aug 24, 2027		U-1556		
	8894988	Aug 24, 2027	DP			
	9073933	Mar 30, 2025	DS			
	9492389	Aug 24, 2027	DP			
	9492391	Aug 24, 2027		U-1556		
	9492392	Aug 24, 2027	DP			
	9492393	Aug 24, 2027		U-1556		
	9522919	Mar 30, 2025	DS DP			
	9763933	Aug 24, 2027	DP			
	9770416	Aug 24, 2027	DP			
	9775808	Aug 24, 2027	DP			
<u>OXYCODONE HYDROCHLORIDE - OXYCONTIN</u>						
N 022272 006	10407434	Mar 30, 2025	DS			
	10696684	Mar 30, 2025	DS			
	11304908	Aug 24, 2027	DP			
	11304909	Aug 24, 2027		U-1556		
	11964056	Aug 24, 2027	DP			
	12060361	Mar 30, 2025	DP	U-1556		
	8808741	Aug 24, 2027		U-1556		
	8894988	Aug 24, 2027	DP			
	9073933	Mar 30, 2025	DS			
	9492389	Aug 24, 2027	DP			
	9492391	Aug 24, 2027		U-1556		
	9492392	Aug 24, 2027	DP			
	9492393	Aug 24, 2027		U-1556		
	9522919	Mar 30, 2025	DS DP			
	9763933	Aug 24, 2027	DP			
	9770416	Aug 24, 2027	DP			
	9775808	Aug 24, 2027	DP			
<u>OXYCODONE HYDROCHLORIDE - OXYCONTIN</u>						
N 022272 007	10407434	Mar 30, 2025	DS			
	10696684	Mar 30, 2025	DS			
	11304908	Aug 24, 2027	DP			
	11304909	Aug 24, 2027		U-1556		
	11964056	Aug 24, 2027	DP			
	12060361	Mar 30, 2025	DP	U-1556		
	8808741	Aug 24, 2027		U-1556		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>OXYCODONE HYDROCHLORIDE - OXYCONTIN</u>						
N 022272	007 8894988	Aug 24, 2027	DP			
	9073933	Mar 30, 2025	DS			
	9492389	Aug 24, 2027	DP			
	9492391	Aug 24, 2027		U-1556		
	9492392	Aug 24, 2027	DP			
	9492393	Aug 24, 2027		U-1556		
	9522919	Mar 30, 2025	DS DP			
	9763933	Aug 24, 2027	DP			
	9770416	Aug 24, 2027	DP			
	9775808	Aug 24, 2027	DP			
<u>OXYCODONE HYDROCHLORIDE - OXAYDO</u>						
N 202080	001 7201920	Mar 16, 2025	DP			
<u>OXYCODONE HYDROCHLORIDE - OXAYDO</u>						
N 202080	002 7201920	Mar 16, 2025	DP			
<u>OXYCODONE HYDROCHLORIDE - ROXYBOND</u>						
N 209777	001 10314788	Aug 12, 2028	DP			
	7955619	Aug 12, 2028	DP			
<u>OXYCODONE HYDROCHLORIDE - ROXYBOND</u>						
N 209777	002 10314788	Aug 12, 2028	DP			
	7955619	Aug 12, 2028	DP			
<u>OXYCODONE HYDROCHLORIDE - ROXYBOND</u>						
N 209777	003 10314788	Aug 12, 2028	DP			
	7955619	Aug 12, 2028	DP			
<u>OXYCODONE HYDROCHLORIDE - ROXYBOND</u>						
N 209777	004 10314788	Aug 12, 2028	DP			
	7955619	Aug 12, 2028	DP			
<u>OXYMETAZOLINE HYDROCHLORIDE - RHOFADÉ</u>						
N 208552	001 10335391	Jun 11, 2035		U-2567		
	10751325	Jun 11, 2035		U-2921		
	11517560	Jun 11, 2035		U-3494		
	7812049	May 02, 2028		U-1959		
	8883838	Dec 01, 2031	DP			
	9974773	Jun 11, 2035		U-2306		
<u>OXYMETAZOLINE HYDROCHLORIDE - UPNEEQ</u>						
N 212520	001 10799481	Dec 16, 2039		U-2849		
	10814001	Dec 16, 2039	DP			
	10898573	Dec 16, 2039	DP			
	10912765	Aug 26, 2031		U-2849		
	10940138	Dec 16, 2039		U-2849		
	11103482	Dec 16, 2039	DP			
	11311515	Dec 16, 2039	DP			
	11324722	Dec 16, 2039		U-2849		
	11541036	Dec 16, 2039	DP			
	11701343	Dec 16, 2039		U-2849		
	8357714	Aug 26, 2031		U-2849		
	9867808	Aug 26, 2031		U-2849		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>OXYMETAZOLINE HYDROCHLORIDE; TETRACAINE HYDROCHLORIDE - KOVANAZE</u>						
N 208032	001	8580282	Apr 02, 2030			DP U-1876
		9308191	Apr 02, 2030			DP U-1876
<u>OXYMORPHONE HYDROCHLORIDE - OPANA ER</u>						
N 021610	001	8808737	Jun 21, 2027			U-3085
		8871779	Nov 22, 2029			DS
<u>OXYMORPHONE HYDROCHLORIDE - OPANA ER</u>						
N 021610	002	8808737	Jun 21, 2027			U-3085
		8871779	Nov 22, 2029			DS
<u>OXYMORPHONE HYDROCHLORIDE - OPANA ER</u>						
N 021610	003	8808737	Jun 21, 2027			U-3085
		8871779	Nov 22, 2029			DS
<u>OXYMORPHONE HYDROCHLORIDE - OPANA ER</u>						
N 021610	004	8808737	Jun 21, 2027			U-3085
		8871779	Nov 22, 2029			DS
<u>OXYMORPHONE HYDROCHLORIDE - OPANA ER</u>						
N 021610	005	8808737	Jun 21, 2027			U-3085
		8871779	Nov 22, 2029			DS
<u>OXYMORPHONE HYDROCHLORIDE - OPANA ER</u>						
N 021610	006	8808737	Jun 21, 2027			U-3085
		8871779	Nov 22, 2029			DS
<u>OXYMORPHONE HYDROCHLORIDE - OPANA ER</u>						
N 021610	007	8808737	Jun 21, 2027			U-3085
		8871779	Nov 22, 2029			DS
<u>OXYMORPHONE HYDROCHLORIDE - OPANA ER</u>						
N 201655	001	7851482	Jul 10, 2029			DS
		8192722	Sep 15, 2025			DP
		8808737	Jun 21, 2027			U-1598
		8871779	Nov 22, 2029			DS
<u>OXYMORPHONE HYDROCHLORIDE - OPANA ER</u>						
N 201655	002	7851482	Jul 10, 2029			DS
		8192722	Sep 15, 2025			DP
		8808737	Jun 21, 2027			U-1598
		8871779	Nov 22, 2029			DS
<u>OXYMORPHONE HYDROCHLORIDE - OPANA ER</u>						
N 201655	003	7851482	Jul 10, 2029			DS
		8192722	Sep 15, 2025			DP
		8808737	Jun 21, 2027			U-1598
		8871779	Nov 22, 2029			DS
<u>OXYMORPHONE HYDROCHLORIDE - OPANA ER</u>						
N 201655	004	7851482	Jul 10, 2029			DS
		8192722	Sep 15, 2025			DP
		8808737	Jun 21, 2027			U-1598
		8871779	Nov 22, 2029			DS

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>OXYMORPHONE HYDROCHLORIDE - OPANA ER</u>						
N 201655	005	7851482	Jul 10, 2029	DS		
		8192722	Sep 15, 2025	DP		
		8808737	Jun 21, 2027		U-1598	
		8871779	Nov 22, 2029	DS		
<u>OXYMORPHONE HYDROCHLORIDE - OPANA ER</u>						
N 201655	006	7851482	Jul 10, 2029	DS		
		8192722	Sep 15, 2025	DP		
		8808737	Jun 21, 2027		U-1598	
		8871779	Nov 22, 2029	DS		
<u>OXYMORPHONE HYDROCHLORIDE - OPANA ER</u>						
N 201655	007	7851482	Jul 10, 2029	DS		
		8192722	Sep 15, 2025	DP		
		8808737	Jun 21, 2027		U-1598	
		8871779	Nov 22, 2029	DS		
<u>OZANIMOD HYDROCHLORIDE - ZEPOSIA</u>						
N 209899	001	10239846	Nov 15, 2030		U-3740	M-309 Aug 30, 2027
		11680050	Sep 30, 2038	DS DP	U-2774	NCE Mar 25, 2025
		11680050	Sep 30, 2038	DS DP	U-3740	
		8481573	May 14, 2029	DS DP	U-2774	
		8796318	May 14, 2029	DS DP		
		9382217	May 14, 2029		U-2774	
<u>OZANIMOD HYDROCHLORIDE - ZEPOSIA</u>						
N 209899	002	10239846	Nov 15, 2030		U-3740	M-309 Aug 30, 2027
		11680050	Sep 30, 2038	DS DP	U-2774	NCE Mar 25, 2025
		11680050	Sep 30, 2038	DS DP	U-3740	
		8481573	May 14, 2029	DS DP	U-2774	
		8796318	May 14, 2029	DS DP		
		9382217	May 14, 2029		U-2774	
<u>OZANIMOD HYDROCHLORIDE - ZEPOSIA</u>						
N 209899	003	10239846	Nov 15, 2030		U-3740	M-309 Aug 30, 2027
		11680050	Sep 30, 2038	DS DP	U-2774	NCE Mar 25, 2025
		11680050	Sep 30, 2038	DS DP	U-3740	
		8481573	May 14, 2029	DS DP	U-2774	
		8796318	May 14, 2029	DS DP		
		9382217	May 14, 2029		U-2774	
<u>OZENOXACIN - XEPI</u>						
N 208945	001	9180200	Jan 29, 2032	DP	U-805	
		9399014	Dec 15, 2029		U-805	
<u>PACLITAXEL - ABRAXANE</u>						
N 021660	001	7758891	Feb 21, 2026		U-1434	
		7758891*PED	Aug 21, 2026			
		7820788*PED	Apr 27, 2025			
		8034375	Aug 13, 2026		U-1290	
		8268348	Feb 21, 2026		U-1290	
		9101543	Feb 21, 2026		U-1434	
		9101543*PED	Aug 21, 2026			
		9393318	Mar 04, 2032		U-1290	

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>PACLITAXEL - ABRAXANE</u>						
N 021660	001	9393318*PED	Sep 04, 2032			
		9511046	Jan 12, 2034	U-1434		
		9511046*PED	Jul 12, 2034			
		9597409	Mar 04, 2032	U-1290		
		9597409*PED	Sep 04, 2032			
<u>PACRITINIB CITRATE - VONJO</u>						
N 208712	001	8153632	Jan 17, 2029	DS DP U-3331	NCE	Feb 28, 2027
		8153632	Jan 17, 2029	DS DP U-3332	ODE-397	Feb 28, 2029
		8980873	Mar 25, 2030	DS DP U-3331		
		8980873	Mar 25, 2030	DS DP U-3332		
		9573964	May 05, 2028	U-3331		
		9573964	May 05, 2028	U-3332		
<u>PAFOLACIANINE SODIUM - CYTALUX</u>						
N 214907	001	10881747	Aug 26, 2033	DS DP U-3291	I-905	Dec 16, 2025
		9061057	Aug 26, 2033	DS DP U-3291	NCE	Nov 29, 2026
		9254341	Oct 04, 2033	DS DP	ODE-390	Nov 29, 2028
		9333270	Aug 26, 2033	DS DP U-3291		
		9341629	Aug 26, 2033	DS DP		
		9789208	Aug 26, 2033	DS DP U-3291		
<u>PALBOCICLIB - IBRANCE</u>						
N 207103	001	10723730	Feb 08, 2034	DS DP		
		RE47739	Mar 05, 2027	DS DP		
<u>PALBOCICLIB - IBRANCE</u>						
N 207103	002	10723730	Feb 08, 2034	DS DP		
		RE47739	Mar 05, 2027	DS DP		
<u>PALBOCICLIB - IBRANCE</u>						
N 207103	003	10723730	Feb 08, 2034	DS DP		
		RE47739	Mar 05, 2027	DS DP		
<u>PALBOCICLIB - IBRANCE</u>						
N 212436	001	10723730	Feb 08, 2034	DS DP		
		11065250	Aug 19, 2036	DP		
		RE47739	Mar 05, 2027	DS DP		
<u>PALBOCICLIB - IBRANCE</u>						
N 212436	002	10723730	Feb 08, 2034	DS DP		
		11065250	Aug 19, 2036	DP		
		RE47739	Mar 05, 2027	DS DP		
<u>PALBOCICLIB - IBRANCE</u>						
N 212436	003	10723730	Feb 08, 2034	DS DP		
		11065250	Aug 19, 2036	DP		
		RE47739	Mar 05, 2027	DS DP		
<u>PALIPERIDONE PALMITATE - INVEGA SUSTENNA</u>						
N 022264	001	9439906	Jan 26, 2031	U-1901		
		9439906	Jan 26, 2031	U-2757		
		9439906	Jan 26, 2031	U-2758		
		9439906	Jan 26, 2031	U-543		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>PALIPERIDONE PALMITATE - INVEGA SUSTENNA</u>						
N 022264 002	9439906	Jan 26, 2031	U-1901			
	9439906	Jan 26, 2031	U-2757			
	9439906	Jan 26, 2031	U-2758			
	9439906	Jan 26, 2031	U-543			
<u>PALIPERIDONE PALMITATE - INVEGA SUSTENNA</u>						
N 022264 003	9439906	Jan 26, 2031	U-1901			
	9439906	Jan 26, 2031	U-2757			
	9439906	Jan 26, 2031	U-2758			
	9439906	Jan 26, 2031	U-543			
<u>PALIPERIDONE PALMITATE - INVEGA SUSTENNA</u>						
N 022264 004	9439906	Jan 26, 2031	U-1901			
	9439906	Jan 26, 2031	U-2757			
	9439906	Jan 26, 2031	U-2758			
	9439906	Jan 26, 2031	U-543			
<u>PALIPERIDONE PALMITATE - INVEGA SUSTENNA</u>						
N 022264 005	9439906	Jan 26, 2031	U-1901			
	9439906	Jan 26, 2031	U-2757			
	9439906	Jan 26, 2031	U-2758			
	9439906	Jan 26, 2031	U-543			
<u>PALIPERIDONE PALMITATE - INVEGA TRINZA</u>						
N 207946 001	10143693	Apr 05, 2036	U-2457			
	10143693	Apr 05, 2036	U-2458			
<u>PALIPERIDONE PALMITATE - INVEGA TRINZA</u>						
N 207946 002	10143693	Apr 05, 2036	U-2457			
	10143693	Apr 05, 2036	U-2458			
<u>PALIPERIDONE PALMITATE - INVEGA TRINZA</u>						
N 207946 003	10143693	Apr 05, 2036	U-2457			
	10143693	Apr 05, 2036	U-2458			
<u>PALIPERIDONE PALMITATE - INVEGA TRINZA</u>						
N 207946 004	10143693	Apr 05, 2036	U-2457			
	10143693	Apr 05, 2036	U-2458			
<u>PALIPERIDONE PALMITATE - INVEGA HAFYERA</u>						
N 207946 005	11304951	May 07, 2041	U-3349			
	11324751	May 07, 2041	U-3359			
	11666697	Nov 24, 2041	U-3626			
<u>PALIPERIDONE PALMITATE - INVEGA HAFYERA</u>						
N 207946 006	11304951	May 07, 2041	U-3349			
	11324751	May 07, 2041	U-3359			
	11666697	Nov 24, 2041	U-3626			
<u>PALIPERIDONE PALMITATE - ERZOFRI</u>						
N 216352 001	11666573	Sep 24, 2039	U-3968			
	11666573	Sep 24, 2039	U-3969			
	12128049	Oct 26, 2038	U-3968			
	12128049	Oct 26, 2038	U-3969			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>PALIPERIDONE PALMITATE - ERZOFRI</u>						
N 216352 002	11666573	Sep 24, 2039	U-3968			
	11666573	Sep 24, 2039	U-3969			
	12128049	Oct 26, 2038	U-3968			
	12128049	Oct 26, 2038	U-3969			
<u>PALIPERIDONE PALMITATE - ERZOFRI</u>						
N 216352 003	11666573	Sep 24, 2039	U-3968			
	11666573	Sep 24, 2039	U-3969			
	12128049	Oct 26, 2038	U-3968			
	12128049	Oct 26, 2038	U-3969			
<u>PALIPERIDONE PALMITATE - ERZOFRI</u>						
N 216352 004	11666573	Sep 24, 2039	U-3968			
	11666573	Sep 24, 2039	U-3969			
	12128049	Oct 26, 2038	U-3968			
	12128049	Oct 26, 2038	U-3969			
<u>PALIPERIDONE PALMITATE - ERZOFRI</u>						
N 216352 005	11666573	Sep 24, 2039	U-3968			
	11666573	Sep 24, 2039	U-3969			
	12128049	Oct 26, 2038	U-3968			
	12128049	Oct 26, 2038	U-3969			
<u>PALIPERIDONE PALMITATE - ERZOFRI</u>						
N 216352 006	11666573	Sep 24, 2039	U-3968			
	11666573	Sep 24, 2039	U-3969			
	12128049	Oct 26, 2038	U-3968			
	12128049	Oct 26, 2038	U-3969			
<u>PALOPEGTERIPARATIDE - YORVIPATH</u>						
N 216490 001	11590207	Sep 28, 2037	U-3982		NP	Aug 09, 2027
	11759504	Sep 28, 2037	DP		ODE-492	Aug 09, 2031
	11857603	Sep 28, 2037	U-3982			
	11890326	Sep 28, 2037	DS DP			
	11918628	Sep 28, 2037	DS DP			
	8906847	Apr 30, 2031	DS DP U-3982			
<u>PALOPEGTERIPARATIDE - YORVIPATH</u>						
N 216490 002	11590207	Sep 28, 2037	U-3982		NP	Aug 09, 2027
	11759504	Sep 28, 2037	DP		ODE-492	Aug 09, 2031
	11857603	Sep 28, 2037	U-3982			
	11890326	Sep 28, 2037	DS DP			
	11918628	Sep 28, 2037	DS DP			
	8906847	Apr 30, 2031	DS DP U-3982			
<u>PALOPEGTERIPARATIDE - YORVIPATH</u>						
N 216490 003	11590207	Sep 28, 2037	U-3982		NP	Aug 09, 2027
	11759504	Sep 28, 2037	DP		ODE-492	Aug 09, 2031
	11857603	Sep 28, 2037	U-3982			
	11890326	Sep 28, 2037	DS DP			
	11918628	Sep 28, 2037	DS DP			
	8906847	Apr 30, 2031	DS DP U-3982			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>PALOVAROTENE - SOHONOS</u>						
N 215559 001	10292954	Aug 31, 2031	U-3676		NCE	Aug 16, 2028
	10864194	Jun 08, 2037	U-3676		ODE-439	Aug 16, 2030
	11622959	Jun 08, 2037	U-3676			
	12023312	Aug 31, 2031	U-3966			
	12138245	Jun 08, 2037	U-4031			
	9314439	Aug 31, 2031	U-3676			
	9789074	Aug 31, 2031	U-3676			
<u>PALOVAROTENE - SOHONOS</u>						
N 215559 002	10292954	Aug 31, 2031	U-3676		NCE	Aug 16, 2028
	10864194	Jun 08, 2037	U-3676		ODE-439	Aug 16, 2030
	11622959	Jun 08, 2037	U-3676			
	12023312	Aug 31, 2031	U-3966			
	12138245	Jun 08, 2037	U-4031			
	9314439	Aug 31, 2031	U-3676			
	9789074	Aug 31, 2031	U-3676			
<u>PALOVAROTENE - SOHONOS</u>						
N 215559 003	10292954	Aug 31, 2031	U-3676		NCE	Aug 16, 2028
	10864194	Jun 08, 2037	U-3676		ODE-439	Aug 16, 2030
	11622959	Jun 08, 2037	U-3676			
	12023312	Aug 31, 2031	U-3966			
	12138245	Jun 08, 2037	U-4031			
	9314439	Aug 31, 2031	U-3676			
	9789074	Aug 31, 2031	U-3676			
<u>PALOVAROTENE - SOHONOS</u>						
N 215559 004	10292954	Aug 31, 2031	U-3676		NCE	Aug 16, 2028
	10864194	Jun 08, 2037	U-3676		ODE-439	Aug 16, 2030
	11622959	Jun 08, 2037	U-3676			
	12023312	Aug 31, 2031	U-3966			
	12138245	Jun 08, 2037	U-4031			
	9314439	Aug 31, 2031	U-3676			
	9789074	Aug 31, 2031	U-3676			
<u>PALOVAROTENE - SOHONOS</u>						
N 215559 005	10292954	Aug 31, 2031	U-3676		NCE	Aug 16, 2028
	10864194	Jun 08, 2037	U-3676		ODE-439	Aug 16, 2030
	11622959	Jun 08, 2037	U-3676			
	12023312	Aug 31, 2031	U-3966			
	12138245	Jun 08, 2037	U-4031			
	9314439	Aug 31, 2031	U-3676			
	9789074	Aug 31, 2031	U-3676			
<u>PANOBINOSTAT LACTATE - FARYDAK</u>						
N 205353 001	7989494	Jan 17, 2028	DS DP			
	8883842	Jun 13, 2028	U-1669			
<u>PANOBINOSTAT LACTATE - FARYDAK</u>						
N 205353 002	7989494	Jan 17, 2028	DS DP			
	8883842	Jun 13, 2028	U-1669			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>PANOBINOSTAT LACTATE - FARYDAK</u>						
N 205353	003 7989494	Jan 17, 2028	DS DP			
	8883842	Jun 13, 2028		U-1669		
<u>PANTOPRAZOLE SODIUM - PROTONIX IV</u>						
N 020988	001				NPP	Aug 12, 2027
<u>PANTOPRAZOLE SODIUM - PROTONIX</u>						
N 022020	001 7544370	Jun 07, 2026	DP			
<u>PAROXETINE MESYLATE - PEXEVA</u>						
N 021299	001 7598271	May 04, 2025	DS			
<u>PAROXETINE MESYLATE - PEXEVA</u>						
N 021299	002 7598271	May 04, 2025	DS			
<u>PAROXETINE MESYLATE - PEXEVA</u>						
N 021299	003 7598271	May 04, 2025	DS			
<u>PAROXETINE MESYLATE - PEXEVA</u>						
N 021299	004 7598271	May 04, 2025	DS			
<u>PAROXETINE MESYLATE - BRISDELLE</u>						
N 204516	001 7598271	May 04, 2025	DS			
	8658663	Apr 06, 2029	DS DP	U-904		
	8946251	Aug 04, 2026	DS DP	U-904		
	9393237	Aug 04, 2026		U-904		
<u>PASIREOTIDE DIASPARTATE - SIGNIFOR</u>						
N 200677	001 7473761	Dec 14, 2026	DS DP			
	8299209	Dec 27, 2025	DS DP			
<u>PASIREOTIDE DIASPARTATE - SIGNIFOR</u>						
N 200677	002 7473761	Dec 14, 2026	DS DP			
	8299209	Dec 27, 2025	DS DP			
<u>PASIREOTIDE DIASPARTATE - SIGNIFOR</u>						
N 200677	003 7473761	Dec 14, 2026	DS DP			
	8299209	Dec 27, 2025	DS DP			
<u>PASIREOTIDE PAMOATE - SIGNIFOR LAR KIT</u>						
N 203255	001 7473761	Dec 14, 2026	DS DP		ODE-268	Jun 29, 2025
	7759308	Oct 25, 2026	DP			
	9351923	May 23, 2028	DP			
<u>PASIREOTIDE PAMOATE - SIGNIFOR LAR KIT</u>						
N 203255	002 7473761	Dec 14, 2026	DS DP		ODE-268	Jun 29, 2025
	7759308	Oct 25, 2026	DP			
	9351923	May 23, 2028	DP			
<u>PASIREOTIDE PAMOATE - SIGNIFOR LAR KIT</u>						
N 203255	003 7473761	Dec 14, 2026	DS DP		ODE-268	Jun 29, 2025
	7759308	Oct 25, 2026	DP			
	9351923	May 23, 2028	DP			
<u>PASIREOTIDE PAMOATE - SIGNIFOR LAR KIT</u>						
N 203255	004 7473761	Dec 14, 2026	DS DP		ODE-268	Jun 29, 2025
	7759308	Oct 25, 2026	DP			
	9351923	May 23, 2028	DP			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>PASIREOTIDE PAMOATE - SIGNIFOR LAR KIT</u>						
N 203255	005 7473761	Dec 14, 2026	DS DP		ODE-268	Jun 29, 2025
	7759308	Oct 25, 2026	DP			
	9351923	May 23, 2028	DP			
<u>PATIOMER SORBITEX CALCIUM - VELTASSA</u>						
N 205739	001 11123363	Oct 08, 2033		U-1766	NPP	Oct 02, 2026
	7556799	Feb 27, 2025		U-1766		
	8147873	Jun 20, 2028	DP			
	8216560	Mar 14, 2027		U-1766		
	8282913	May 29, 2027	DP			
	8337824	May 29, 2030	DS	U-1766		
	9492476	Oct 08, 2033		U-1766		
	9925212	Oct 08, 2033		U-1766		
<u>PATIOMER SORBITEX CALCIUM - VELTASSA</u>						
N 205739	002 11123363	Oct 08, 2033		U-1766	NPP	Oct 02, 2026
	7556799	Feb 27, 2025		U-1766		
	8147873	Jun 20, 2028	DP			
	8216560	Mar 14, 2027		U-1766		
	8282913	May 29, 2027	DP			
	8337824	May 29, 2030	DS	U-1766		
	9492476	Oct 08, 2033		U-1766		
	9925212	Oct 08, 2033		U-1766		
<u>PATIOMER SORBITEX CALCIUM - VELTASSA</u>						
N 205739	003 11123363	Oct 08, 2033		U-1766	NPP	Oct 02, 2026
	7556799	Feb 27, 2025		U-1766		
	8147873	Jun 20, 2028	DP			
	8216560	Mar 14, 2027		U-1766		
	8282913	May 29, 2027	DP			
	8337824	May 29, 2030	DS	U-1766		
	9492476	Oct 08, 2033		U-1766		
	9925212	Oct 08, 2033		U-1766		
<u>PATIOMER SORBITEX CALCIUM - VELTASSA</u>						
N 205739	004 11123363	Oct 08, 2033		U-1766	NS	Oct 02, 2026
	7556799	Feb 27, 2025		U-1766		
	8147873	Jun 20, 2028	DP			
	8216560	Mar 14, 2027		U-1766		
	8282913	May 29, 2027	DP			
	8337824	May 29, 2030	DS	U-1766		
	9492476	Oct 08, 2033		U-1766		
	9925212	Oct 08, 2033		U-1766		
<u>PATISIRAN SODIUM - ONPATTRO</u>						
N 210922	001 10240152	Oct 20, 2029	DS DP	U-2378	M-270	Jan 13, 2026
	11079379	Aug 27, 2035	DS DP	U-2378	ODE-197	Aug 10, 2025
	11141378	Apr 15, 2029	DP			
	8058069	Apr 15, 2029	DP			
	8158601	Nov 10, 2030	DP	U-2378		
	8168775	Oct 20, 2029	DS DP	U-2378		
	8334373	May 27, 2025	DS DP			
	8492359	Apr 15, 2029	DP			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>PATISIRAN SODIUM - ONPATRO</u>						
N 210922	001 8642076	Oct 03, 2027	DP			
	8741866	Oct 20, 2029	U-2378			
	8802644	Oct 21, 2030	DP U-2378			
	8822668	Apr 15, 2029	DP U-2378			
	9234196	Oct 20, 2029	DP U-2378			
	9364435	Apr 15, 2029	DP U-2378			
<u>PEGCETACOPLAN - EMPAVELI</u>						
N 215014	001 10035822	Nov 15, 2033	DS		M-288	Feb 08, 2026
	10125171	Aug 02, 2033	DS		NCE	May 14, 2026
	10875893	Nov 15, 2033	DS U-3124		ODE-351	May 14, 2028
	11040107	Apr 09, 2038	DP U-3172			
	11040107	Apr 09, 2038	DP U-3173			
	11040107	Apr 09, 2038	DP U-3174			
	11292815	Nov 15, 2033	DS DP U-3124			
	11292815	Nov 15, 2033	DS DP U-3354			
	11661441	Jan 13, 2033	DS U-3124			
	11844841	Dec 09, 2038	DP U-3172			
	11844841	Dec 09, 2038	DP U-3173			
	11844841	Dec 09, 2038	DP U-3174			
	7888323	Dec 04, 2027	DS			
	7989589	Dec 04, 2027	DS			
	9169307	Nov 18, 2027	DS U-3123			
<u>PEGCETACOPLAN - SYFOVRE</u>						
N 217171	001 10035822	Nov 15, 2033	DS		NCE	May 14, 2026
	10125171	Aug 02, 2033	DS		NP	Feb 22, 2026
	10875893	Nov 15, 2033	DS U-3540			
	11292815	Nov 15, 2033	DS DP U-3540			
	11661441	Jan 13, 2033	DS U-3540			
	11903994	Feb 22, 2037	U-3826			
	11903994	Feb 22, 2037	U-3827			
	11903994	Feb 22, 2037	U-3828			
	7888323	Dec 04, 2027	DS			
	7989589	Dec 04, 2027	DS			
	8168584	Apr 07, 2027	U-3540			
	8168584	Apr 07, 2027	U-3542			
	9056076	Oct 25, 2026	U-3540			
	9169307	Nov 18, 2027	DS U-3541			
<u>PEGINESATIDE ACETATE - OMONTYS PRESERVATIVE FREE</u>						
N 202799	001 7550433	Jun 02, 2026	U-1238			
	7919461	Jun 02, 2026	U-1238			
<u>PEGINESATIDE ACETATE - OMONTYS PRESERVATIVE FREE</u>						
N 202799	002 7550433	Jun 02, 2026	U-1238			
	7919461	Jun 02, 2026	U-1238			
<u>PEGINESATIDE ACETATE - OMONTYS PRESERVATIVE FREE</u>						
N 202799	003 7550433	Jun 02, 2026	U-1238			
	7919461	Jun 02, 2026	U-1238			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>PEGINESATIDE ACETATE - OMONTYS PRESERVATIVE FREE</u>						
N 202799 004	7550433	Jun 02, 2026	U-1238			
	7919461	Jun 02, 2026	U-1238			
<u>PEGINESATIDE ACETATE - OMONTYS PRESERVATIVE FREE</u>						
N 202799 005	7550433	Jun 02, 2026	U-1238			
	7919461	Jun 02, 2026	U-1238			
<u>PEGINESATIDE ACETATE - OMONTYS PRESERVATIVE FREE</u>						
N 202799 006	7550433	Jun 02, 2026	U-1238			
	7919461	Jun 02, 2026	U-1238			
<u>PEGINESATIDE ACETATE - OMONTYS</u>						
N 202799 007	7550433	Jun 02, 2026	U-1238			
	7919461	Jun 02, 2026	U-1238			
<u>PEGINESATIDE ACETATE - OMONTYS</u>						
N 202799 008	7550433	Jun 02, 2026	U-1238			
	7919461	Jun 02, 2026	U-1238			
<u>PEGULICIANINE ACETATE - LUMISIGHT</u>						
N 214511 001	10285759	Dec 08, 2031	U-3890		NCE	Apr 17, 2029
	9032965	Dec 08, 2031	U-3890			
	9155471	Oct 12, 2031	U-3890			
	9532835	Dec 08, 2031	U-3890			
	9763577	Sep 14, 2034	DS DP U-3890			
<u>PEMETREXED - PEMFEXY</u>						
N 209472 001	11793813	Feb 19, 2036	DP			
	9604990	Oct 28, 2035	DS			
<u>PEMETREXED DISODIUM - PEMETREXED</u>						
N 215179 001	11147817	Mar 26, 2035	DP			
<u>PEMETREXED DISODIUM - PEMETREXED</u>						
N 215179 002	11147817	Mar 26, 2035	DP			
<u>PEMETREXED DISODIUM - PEMETREXED</u>						
N 215179 003	11147817	Mar 26, 2035	DP			
<u>PEMIGATINIB - PEMAZYRE</u>						
N 213736 001	10131667	Jun 12, 2033	U-2809		I-899	Aug 26, 2025
	11466004	May 03, 2039	U-3464		NCE	Apr 17, 2025
	11466004	May 03, 2039	U-3465		ODE-292	Apr 17, 2027
	11466004	May 03, 2039	U-3466		ODE-404	Aug 26, 2029
	11628162	Aug 30, 2040	U-3568			
	11628162	Aug 30, 2040	U-3569			
	11628162	Aug 30, 2040	U-3570			
	11628162	Aug 30, 2040	U-3571			
	9611267	Jan 30, 2035	DS DP			
<u>PEMIGATINIB - PEMAZYRE</u>						
N 213736 002	10131667	Jun 12, 2033	U-2809		I-899	Aug 26, 2025
	11466004	May 03, 2039	U-3464		NCE	Apr 17, 2025
	11466004	May 03, 2039	U-3465		ODE-292	Apr 17, 2027
	11466004	May 03, 2039	U-3466		ODE-404	Aug 26, 2029
	11628162	Aug 30, 2040	U-3568			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>PEMIGATINIB - PEMAZYRE</u>						
N 213736	002	11628162	Aug 30, 2040	U-3569		
		11628162	Aug 30, 2040	U-3570		
		11628162	Aug 30, 2040	U-3571		
		9611267	Jan 30, 2035	DS DP		
<u>PEMIGATINIB - PEMAZYRE</u>						
N 213736	003	10131667	Jun 12, 2033	U-2809	I-899	Aug 26, 2025
		11466004	May 03, 2039	U-3464	NCE	Apr 17, 2025
		11466004	May 03, 2039	U-3465	ODE-292	Apr 17, 2027
		11466004	May 03, 2039	U-3466	ODE-404	Aug 26, 2029
		11628162	Aug 30, 2040	U-3568		
		11628162	Aug 30, 2040	U-3569		
		11628162	Aug 30, 2040	U-3570		
		11628162	Aug 30, 2040	U-3571		
		9611267	Jan 30, 2035	DS DP		
<u>PERAMIVIR - RAPIVAB</u>						
N 206426	001	10391075	Feb 12, 2027	U-2622		
		10391075	Feb 12, 2027	U-3069		
		8778997	May 07, 2027	U-1627		
		8778997	May 07, 2027	U-2622		
		8778997	May 07, 2027	U-3069		
<u>PERAMPANEL - FYCOMPA</u>						
N 202834	001	6949571	May 23, 2025	DS DP U-106		
		6949571	May 23, 2025	DS DP U-2088		
		6949571	May 23, 2025	DS DP U-2089		
		6949571	May 23, 2025	DS DP U-2428		
		6949571	May 23, 2025	DS DP U-2429		
		8772497	Jul 01, 2026	DS		
<u>PERAMPANEL - FYCOMPA</u>						
N 202834	002	6949571	May 23, 2025	DS DP U-106		
		6949571	May 23, 2025	DS DP U-2088		
		6949571	May 23, 2025	DS DP U-2089		
		6949571	May 23, 2025	DS DP U-2428		
		6949571	May 23, 2025	DS DP U-2429		
		8772497	Jul 01, 2026	DS		
<u>PERAMPANEL - FYCOMPA</u>						
N 202834	003	6949571	May 23, 2025	DS DP U-106		
		6949571	May 23, 2025	DS DP U-2088		
		6949571	May 23, 2025	DS DP U-2089		
		6949571	May 23, 2025	DS DP U-2428		
		6949571	May 23, 2025	DS DP U-2429		
		8772497	Jul 01, 2026	DS		
<u>PERAMPANEL - FYCOMPA</u>						
N 202834	004	6949571	May 23, 2025	DS DP U-106		
		6949571	May 23, 2025	DS DP U-2088		
		6949571	May 23, 2025	DS DP U-2089		
		6949571	May 23, 2025	DS DP U-2428		
		6949571	May 23, 2025	DS DP U-2429		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>PERAMPANEL - FYCOMPA</u>						
N 202834	004	8772497	Jul 01, 2026	DS		
<u>PERAMPANEL - FYCOMPA</u>						
N 202834	005	6949571	May 23, 2025	DS DP U-106		
		6949571	May 23, 2025	DS DP U-2088		
		6949571	May 23, 2025	DS DP U-2089		
		6949571	May 23, 2025	DS DP U-2428		
		6949571	May 23, 2025	DS DP U-2429		
		8772497	Jul 01, 2026	DS		
<u>PERAMPANEL - FYCOMPA</u>						
N 202834	006	6949571	May 23, 2025	DS DP U-106		
		6949571	May 23, 2025	DS DP U-2088		
		6949571	May 23, 2025	DS DP U-2089		
		6949571	May 23, 2025	DS DP U-2428		
		6949571	May 23, 2025	DS DP U-2429		
		8772497	Jul 01, 2026	DS		
<u>PERAMPANEL - FYCOMPA</u>						
N 208277	001	6949571	May 23, 2025	DS DP U-106		
		6949571	May 23, 2025	DS DP U-2088		
		6949571	May 23, 2025	DS DP U-2089		
		6949571	May 23, 2025	DS DP U-2428		
		6949571	May 23, 2025	DS DP U-2429		
		8772497	Jul 01, 2026	DS		
<u>PERFLUOROHEXYLOCTANE - MIEBO</u>						
N 216675	001	10058615	Sep 12, 2033	U-1900	NCE	May 18, 2028
		10369117	Sep 12, 2033	U-1900		
		10449164	Sep 12, 2033	U-1900		
		10507132	Jun 21, 2037	U-1900		
		10576154	Sep 12, 2033	U-1900		
		11357738	Sep 29, 2036	DP		
<u>PERFLUTREN - DEFINITY</u>						
N 021064	001	10583207	Dec 28, 2035	U-665		
		10583208	Mar 16, 2037	U-665		
		10588988	May 04, 2037	U-665		
		11266750	Mar 16, 2037	U-665		
		11529431	Mar 16, 2037	U-665		
		11857646	Mar 16, 2037	U-665		
		11925695	Mar 16, 2037	U-665		
		12161730	Mar 16, 2037	U-665		
		9789210	Mar 16, 2037	U-665		
<u>PERFLUTREN - DEFINITY RT</u>						
N 021064	002	10022460	Dec 28, 2035	DS DP		
		10583207	Dec 28, 2035	U-665		
		10583208	Mar 16, 2037	U-665		
		10588988	May 04, 2037	U-665		
		11266750	Mar 16, 2037	U-665		
		11395856	Dec 28, 2035	DS DP U-665		
		11529431	Mar 16, 2037	U-665		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>PERFLUTREN - DEFINITY RT</u>						
N 021064	002	11857646	Mar 16, 2037	U-665		
		11925695	Mar 16, 2037	U-665		
		12161730	Mar 16, 2037	U-665		
		9789210	Mar 16, 2037	U-665		
<u>PEXIDARTINIB HYDROCHLORIDE - TURALIO</u>						
N 211810	001	10189833	May 05, 2036	U-2606	ODE-250	Aug 02, 2026
		10435404	Jul 24, 2038	DP		
		10730876	May 05, 2036	DS		
		10941142	Jul 24, 2038	DP		
		10961240	Jul 24, 2038	U-2606		
		7893075	May 04, 2033	DS		
		8404700	Nov 21, 2027	DS		
		8461169	Apr 19, 2028	U-2606		
		8722702	Nov 21, 2027	DS		
		9169250	Nov 21, 2027	DS		
		9358235	Jun 08, 2033	U-2606		
		9802932	May 05, 2036	DS		
<u>PEXIDARTINIB HYDROCHLORIDE - TURALIO</u>						
N 211810	002	10189833	May 05, 2036	U-2606	ODE*	Aug 02, 2026
		10435404	Jul 24, 2038	DP		
		10730876	May 05, 2036	DS		
		10941142	Jul 24, 2038	DP		
		10961240	Jul 24, 2038	U-2606		
		7893075	May 04, 2033	DS		
		8404700	Nov 21, 2027	DS		
		8461169	Apr 19, 2028	U-2606		
		8722702	Nov 21, 2027	DS		
		9169250	Nov 21, 2027	DS		
		9358235	Jun 08, 2033	U-2606		
		9802932	May 05, 2036	DS		
<u>PHENOBARBITAL SODIUM - SEZABY</u>						
N 215910	001	11857683	Apr 07, 2042	DP U-3779	ODE-414	Nov 17, 2029
<u>PHENTERMINE HYDROCHLORIDE - SUPRENZA</u>						
N 202088	001	8440170	Mar 14, 2029	DP		
<u>PHENTERMINE HYDROCHLORIDE - SUPRENZA</u>						
N 202088	002	8440170	Mar 14, 2029	DP		
<u>PHENTERMINE HYDROCHLORIDE - SUPRENZA</u>						
N 202088	003	8440170	Mar 14, 2029	DP		
<u>PHENTERMINE HYDROCHLORIDE; TOPIRAMATE - QSYMIA</u>						
N 022580	001	8580298	May 15, 2029	DP	M-308	Sep 13, 2027
		8580299	Jun 14, 2029	U-3399	NPP	Jun 24, 2025
		8895057	Jun 09, 2028	U-3398		
		8895058	Jun 09, 2028	DP		
		9011905	Jun 09, 2028	DP		
		9011906	Jun 09, 2028	U-3398		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>PHENTERMINE HYDROCHLORIDE; TOPIRAMATE - QSYMIA</u>						
N 022580 002	8580298	May 15, 2029	DP		M-308	Sep 13, 2027
	8580299	Jun 14, 2029	U-3399		NPP	Jun 24, 2025
	8895057	Jun 09, 2028	U-3398			
	8895058	Jun 09, 2028	DP			
	9011905	Jun 09, 2028	DP			
	9011906	Jun 09, 2028	U-3398			
<u>PHENTERMINE HYDROCHLORIDE; TOPIRAMATE - OSYMIA</u>						
N 022580 003	8580298	May 15, 2029	DP		M-308	Sep 13, 2027
	8580299	Jun 14, 2029	U-3399		NPP	Jun 24, 2025
	8895057	Jun 09, 2028	U-3398			
	8895058	Jun 09, 2028	DP			
	9011905	Jun 09, 2028	DP			
	9011906	Jun 09, 2028	U-3398			
<u>PHENTERMINE HYDROCHLORIDE; TOPIRAMATE - OSYMIA</u>						
N 022580 004	8580298	May 15, 2029	DP		M-308	Sep 13, 2027
	8580299	Jun 14, 2029	U-3399		NPP	Jun 24, 2025
	8895057	Jun 09, 2028	U-3398			
	8895058	Jun 09, 2028	DP			
	9011905	Jun 09, 2028	DP			
	9011906	Jun 09, 2028	U-3398			
<u>PHENTOLAMINE MESYLATE - RYZUMVI</u>						
N 217064 001	10278918	Jan 31, 2034	DP		NP	Sep 25, 2026
	10772829	Jan 31, 2034	DP			
	11090261	Jan 31, 2034	DP			
	11400077	Oct 25, 2039	U-3804			
	11844858	Jan 31, 2034	DP			
	9795560	Jan 31, 2034	DP			
<u>PHENYLEPHRINE HYDROCHLORIDE - PHENYLEPHRINE HYDROCHLORIDE</u>						
N 203510 001	8859623	Nov 14, 2033	U-1594			
<u>PHENYLEPHRINE HYDROCHLORIDE - PHENYLEPHRINE HYDROCHLORIDE</u>						
N 203510 002	8859623	Nov 14, 2033	U-1594			
<u>PHENYLEPHRINE HYDROCHLORIDE - IMPHENTIV</u>						
N 203826 004	11213480	Sep 26, 2036	DP			
	11471400	Aug 05, 2036	DP			
<u>PHENYLEPHRINE HYDROCHLORIDE - IMPHENTIV</u>						
N 203826 005	11213480	Sep 26, 2036	DP			
	11471400	Aug 05, 2036	DP			
<u>PHENYLEPHRINE HYDROCHLORIDE; TROPICAMIDE - MYDCOMBI</u>						
N 215352 001	10839960	Jul 15, 2031	U-3685		NP	May 05, 2026
	11398306	Jul 15, 2031	U-3685			
	11839487	Jul 15, 2031	U-3685			
<u>PIFLUFOLASTAT F-18 - PYLARIFY</u>						
N 214793 001	10947197	Jun 09, 2037	DS DP U-3130		NCE	May 26, 2026
	11851407	Jun 09, 2037	DS DP U-3130			
	12070513	Jul 31, 2029	DS DP U-3130			
	8487129	Nov 07, 2027	DS DP			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>PIFLUFOLASTAT F-18 - PYLARIFY</u>						
N 214793	001 8778305	Sep 21, 2030	DS DP U-3130			
	9861713	Jul 31, 2029	DS DP U-3130			
<u>PILOCARPINE HYDROCHLORIDE - VUITY</u>						
N 214028	001 10610518	Apr 24, 2039	U-3252		D-187	Mar 28, 2026
	10610518	Apr 24, 2039	U-3561			
	10610518	Apr 24, 2039	U-3562			
	11285134	Apr 24, 2039	U-3252			
	11285134	Apr 24, 2039	U-3561			
	11285134	Apr 24, 2039	U-3562			
<u>PILOCARPINE HYDROCHLORIDE - QLOSI</u>						
N 217836	001 10639297	Aug 18, 2037	DP U-3741		NP	Oct 17, 2026
	11129812	Aug 18, 2037	U-3741			
	11974986	Aug 18, 2037	U-3741			
	9867810	Aug 18, 2037	DP U-3741			
<u>PIMAVANSERIN TARTRATE - NUPLAZID</u>						
N 207318	001 7601740	Apr 29, 2030	DS DP			
	7659285	Aug 24, 2026	U-1844			
	7732615	Jun 03, 2028	DS DP			
	7923564	Sep 26, 2025	DS DP			
<u>PIMAVANSERIN TARTRATE - NUPLAZID</u>						
N 207318	002 10517860	Mar 23, 2037	U-1974			
	10953000	Mar 23, 2037	U-1974			
	7601740	Apr 29, 2030	DS DP			
	7659285	Aug 24, 2026	U-1844			
	7732615	Jun 03, 2028	DS DP			
	7923564	Sep 26, 2025	DS DP			
<u>PIMAVANSERIN TARTRATE - NUPLAZID</u>						
N 210793	001 10449185	Aug 27, 2038	DP			
	10646480	Aug 27, 2038	DP			
	10849891	Aug 27, 2038	DP U-1974			
	11452721	Aug 27, 2038	DP			
	7601740	Apr 29, 2030	DS DP			
	7659285	Aug 24, 2026	U-1844			
	7732615	Jun 03, 2028	DS DP			
	7923564	Sep 26, 2025	DS DP			
<u>PIRFENIDONE - ESBRIET</u>						
N 022535	001 7566729	Apr 22, 2029	U-1600			
	7635707	Apr 22, 2029	U-1609			
	7696236	Dec 18, 2027	U-1601			
	7767225	Sep 22, 2026	DP U-1602			
	7767700	Dec 18, 2027	U-1601			
	7816383	Jan 08, 2030	U-1603			
	7910610	Jan 08, 2030	U-1604			
	7988994	Sep 22, 2026	DP U-1602			
	8013002	Jan 08, 2030	U-1603			
	8084475	Jan 08, 2030	U-1605			
	8318780	Jan 08, 2030	U-1606			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>PIRFENIDONE - ESBRIET</u>						
N 022535	001 8383150	May 10, 2028	DP U-2361			
	8420674	Dec 18, 2027	DP U-1608			
	8592462	Apr 22, 2029	U-1609			
	8609701	Apr 22, 2029	U-1610			
	8648098	Jan 08, 2030	U-1611			
	8753679	Sep 22, 2026	DP U-1602			
	8754109	Jan 08, 2030	U-1612			
	8778947	Aug 30, 2033	U-1613			
<u>PIRFENIDONE - ESBRIET</u>						
N 208780	001 10188637	Mar 28, 2037	DP			
	7566729	Apr 22, 2029	U-2077			
	7566729	Apr 22, 2029	U-2078			
	7635707	Apr 22, 2029	U-2072			
	7635707	Apr 22, 2029	U-2073			
	7635707	Apr 22, 2029	U-2074			
	7635707	Apr 22, 2029	U-2075			
	7635707	Apr 22, 2029	U-2076			
	7635707	Apr 22, 2029	U-2083			
	7767700	Dec 18, 2027	U-2080			
	7816383	Jan 08, 2030	U-2042			
	7816383	Jan 08, 2030	U-2050			
	7910610	Jan 08, 2030	U-2048			
	7910610	Jan 08, 2030	U-2049			
	8013002	Jan 08, 2030	U-2047			
	8013002	Jan 08, 2030	U-2082			
	8084475	Jan 08, 2030	U-2052			
	8084475	Jan 08, 2030	U-2054			
	8318780	Jan 08, 2030	U-2046			
	8318780	Jan 08, 2030	U-2081			
	8383150	May 10, 2028	DP U-2361			
	8420674	Dec 18, 2027	U-2079			
	8592462	Apr 22, 2029	U-2055			
	8592462	Apr 22, 2029	U-2056			
	8592462	Apr 22, 2029	U-2057			
	8592462	Apr 22, 2029	U-2058			
	8592462	Apr 22, 2029	U-2059			
	8592462	Apr 22, 2029	U-2060			
	8592462	Apr 22, 2029	U-2061			
	8592462	Apr 22, 2029	U-2062			
	8592462	Apr 22, 2029	U-2063			
	8609701	Apr 22, 2029	U-2064			
	8609701	Apr 22, 2029	U-2065			
	8609701	Apr 22, 2029	U-2066			
	8609701	Apr 22, 2029	U-2067			
	8609701	Apr 22, 2029	U-2068			
	8609701	Apr 22, 2029	U-2069			
	8609701	Apr 22, 2029	U-2070			
	8648098	Jan 08, 2030	U-2051			
	8648098	Jan 08, 2030	U-2052			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>PIRFENIDONE - ESBRIET</u>						
N 208780 001	8754109	Jan 08, 2030	U-2053			
	8778947	Aug 30, 2033	U-2044			
	8778947	Aug 30, 2033	U-2045			
<u>PIRFENIDONE - ESBRIET</u>						
N 208780 002	10188637	Mar 28, 2037	DP			
	7566729	Apr 22, 2029	U-2269			
	7566729	Apr 22, 2029	U-2270			
	7635707	Apr 22, 2029	U-2072			
	7635707	Apr 22, 2029	U-2073			
	7635707	Apr 22, 2029	U-2074			
	7635707	Apr 22, 2029	U-2075			
	7635707	Apr 22, 2029	U-2076			
	7635707	Apr 22, 2029	U-2083			
	7767700	Dec 18, 2027	U-2080			
	7816383	Jan 08, 2030	U-2042			
	7816383	Jan 08, 2030	U-2050			
	7910610	Jan 08, 2030	U-2048			
	7910610	Jan 08, 2030	U-2049			
	8013002	Jan 08, 2030	U-2047			
	8013002	Jan 08, 2030	U-2082			
	8084475	Jan 08, 2030	U-2054			
	8084475	Jan 08, 2030	U-2268			
	8318780	Jan 08, 2030	U-2046			
	8318780	Jan 08, 2030	U-2081			
	8383150	May 10, 2028	DP U-2361			
	8420674	Dec 18, 2027	U-2079			
	8592462	Apr 22, 2029	U-2055			
	8592462	Apr 22, 2029	U-2056			
	8592462	Apr 22, 2029	U-2057			
	8592462	Apr 22, 2029	U-2058			
	8592462	Apr 22, 2029	U-2059			
	8592462	Apr 22, 2029	U-2060			
	8592462	Apr 22, 2029	U-2061			
	8592462	Apr 22, 2029	U-2062			
	8592462	Apr 22, 2029	U-2063			
	8609701	Apr 22, 2029	U-2064			
	8609701	Apr 22, 2029	U-2065			
	8609701	Apr 22, 2029	U-2066			
	8609701	Apr 22, 2029	U-2067			
	8609701	Apr 22, 2029	U-2068			
	8609701	Apr 22, 2029	U-2069			
	8609701	Apr 22, 2029	U-2070			
	8648098	Jan 08, 2030	U-2051			
	8648098	Jan 08, 2030	U-2052			
	8754109	Jan 08, 2030	U-2053			
	8778947	Aug 30, 2033	U-2044			
	8778947	Aug 30, 2033	U-2045			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>PIRFENIDONE - ESBRIET</u>						
N 208780	003 10188637	Mar 28, 2037	DP			
	7566729	Apr 22, 2029		U-2077		
	7566729	Apr 22, 2029		U-2078		
	7635707	Apr 22, 2029		U-2072		
	7635707	Apr 22, 2029		U-2073		
	7635707	Apr 22, 2029		U-2074		
	7635707	Apr 22, 2029		U-2075		
	7635707	Apr 22, 2029		U-2076		
	7635707	Apr 22, 2029		U-2083		
	7767700	Dec 18, 2027		U-2080		
	7816383	Jan 08, 2030		U-2042		
	7816383	Jan 08, 2030		U-2050		
	7910610	Jan 08, 2030		U-2048		
	7910610	Jan 08, 2030		U-2049		
	8013002	Jan 08, 2030		U-2047		
	8013002	Jan 08, 2030		U-2082		
	8084475	Jan 08, 2030		U-2052		
	8084475	Jan 08, 2030		U-2054		
	8318780	Jan 08, 2030		U-2046		
	8318780	Jan 08, 2030		U-2081		
	8383150	May 10, 2028	DP	U-2361		
	8420674	Dec 18, 2027		U-2079		
	8592462	Apr 22, 2029		U-2055		
	8592462	Apr 22, 2029		U-2056		
	8592462	Apr 22, 2029		U-2057		
	8592462	Apr 22, 2029		U-2058		
	8592462	Apr 22, 2029		U-2059		
	8592462	Apr 22, 2029		U-2060		
	8592462	Apr 22, 2029		U-2061		
	8592462	Apr 22, 2029		U-2062		
	8592462	Apr 22, 2029		U-2063		
	8609701	Apr 22, 2029		U-2064		
	8609701	Apr 22, 2029		U-2065		
	8609701	Apr 22, 2029		U-2066		
	8609701	Apr 22, 2029		U-2067		
	8609701	Apr 22, 2029		U-2068		
	8609701	Apr 22, 2029		U-2069		
	8609701	Apr 22, 2029		U-2070		
	8648098	Jan 08, 2030		U-2051		
	8648098	Jan 08, 2030		U-2052		
	8754109	Jan 08, 2030		U-2053		
	8778947	Aug 30, 2033		U-2044		
	8778947	Aug 30, 2033		U-2045		
<u>PIRTOBRUTINIB - JAYPIRCA</u>						
N 216059	001 10342780	Dec 16, 2036	DS DP		NCE	Jan 27, 2028
	10464905	Dec 16, 2036		U-3518	ODE-424	Jan 27, 2030
	10464905	Dec 16, 2036		U-3761	ODE-451	Dec 01, 2030
	10695323	Dec 16, 2036	DS DP	U-3518		
	10695323	Dec 16, 2036	DS DP	U-3761		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>PIRTOBRUTINIB - JAYPIRCA</u>						
N 216059	001 10918622	Dec 16, 2036	U-3518			
	10918622	Dec 16, 2036	U-3761			
	12109193	Sep 14, 2041	DP			
<u>PIRTOBRUTINIB - JAYPIRCA</u>						
N 216059	002 10342780	Dec 16, 2036	DS DP		NCE	Jan 27, 2028
	10464905	Dec 16, 2036	U-3518		ODE-424	Jan 27, 2030
	10464905	Dec 16, 2036	U-3761		ODE-451	Dec 01, 2030
	10695323	Dec 16, 2036	DS DP U-3518			
	10695323	Dec 16, 2036	DS DP U-3761			
	10918622	Dec 16, 2036	U-3518			
	10918622	Dec 16, 2036	U-3761			
	12109193	Sep 14, 2041	DP			
<u>PITAVASTATIN MAGNESIUM - ZYPITAMAG</u>						
N 208379	001 8829186	Jan 19, 2031	DS DP			
<u>PITAVASTATIN MAGNESIUM - ZYPITAMAG</u>						
N 208379	002 8829186	Jan 19, 2031	DS DP			
<u>PITAVASTATIN MAGNESIUM - ZYPITAMAG</u>						
N 208379	003 8829186	Jan 19, 2031	DS DP			
<u>PITOLISANT HYDROCHLORIDE - WAKIX</u>						
N 211150	001 8207197	Mar 07, 2030	DS DP		NPP	Jun 21, 2027
	8354430	Feb 06, 2026	U-1101		ODE-255	Aug 14, 2026
	8354430	Feb 06, 2026	U-1102		ODE-331	Oct 13, 2027
	8486947	Sep 26, 2029	U-1101		ODE-489	Jun 21, 2031
	8486947	Sep 26, 2029	U-1102			
<u>PITOLISANT HYDROCHLORIDE - WAKIX</u>						
N 211150	002 8207197	Mar 07, 2030	DS DP		NPP	Jun 21, 2027
	8354430	Feb 06, 2026	U-1101		ODE-255	Aug 14, 2026
	8354430	Feb 06, 2026	U-1102		ODE-331	Oct 13, 2027
	8486947	Sep 26, 2029	U-1101		ODE-489	Jun 21, 2031
	8486947	Sep 26, 2029	U-1102			
<u>PLAZOMICIN SULFATE - ZEMDRI</u>						
N 210303	001 8383596	Jun 02, 2031	DS U-2328		NCE	Jun 25, 2023
	8822424	Nov 21, 2028	DP		GAIN	Jun 25, 2028
	9266919	Nov 21, 2028	U-2328			
	9688711	Nov 21, 2028	DS U-2328			
<u>PLECANATIDE - TRULANCE</u>						
N 208745	001 10011637	Jun 05, 2034	DS			
	11142549	Jun 05, 2034	DP			
	11319346	Mar 01, 2032	DP			
	11834521	Jun 05, 2034	DP			
	12146003	Jun 05, 2034	DP			
	7041786	Jan 30, 2028	DS			
	9610321	Sep 15, 2031	U-1999			
	9610321	Sep 15, 2031	U-2230			
	9616097	Aug 20, 2032	DP			
	9919024	Sep 15, 2031	U-1999			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>PLECANATIDE - TRULANCE</u>						
N 208745	001 9919024	Sep 15, 2031	U-2230			
	9925231	Sep 15, 2031	DP			
<u>POLIDOCANOL - VARITHENA</u>						
N 205098	001 7814943	Nov 19, 2027	DP U-1461			
	9480652	May 12, 2032	DP			
<u>POMALIDOMIDE - POMALYST</u>						
N 204026	001 10555939	May 19, 2030	DP		ODE-296	May 14, 2027
	10555939*PED	Nov 19, 2030			ODE-297	May 14, 2027
	8198262	Jun 17, 2025	U-1360		PED	Nov 14, 2027
	8198262	Jun 17, 2025	U-2254		PED	Nov 14, 2027
	8198262*PED	Dec 17, 2025				
	8828427	Jun 21, 2031	DS DP			
	8828427*PED	Dec 21, 2031				
	9993467	May 19, 2030	DP			
	9993467*PED	Nov 19, 2030				
<u>POMALIDOMIDE - POMALYST</u>						
N 204026	002 10555939	May 19, 2030	DP		ODE-296	May 14, 2027
	10555939*PED	Nov 19, 2030			ODE-297	May 14, 2027
	8198262	Jun 17, 2025	U-1360		PED	Nov 14, 2027
	8198262	Jun 17, 2025	U-2254		PED	Nov 14, 2027
	8198262*PED	Dec 17, 2025				
	8828427	Jun 21, 2031	DS DP			
	8828427*PED	Dec 21, 2031				
	9993467	May 19, 2030	DP			
	9993467*PED	Nov 19, 2030				
<u>POMALIDOMIDE - POMALYST</u>						
N 204026	003 10555939	May 19, 2030	DP		ODE-296	May 14, 2027
	10555939*PED	Nov 19, 2030			ODE-297	May 14, 2027
	8198262	Jun 17, 2025	U-1360		PED	Nov 14, 2027
	8198262	Jun 17, 2025	U-2254		PED	Nov 14, 2027
	8198262*PED	Dec 17, 2025				
	8828427	Jun 21, 2031	DS DP			
	8828427*PED	Dec 21, 2031				
	9993467	May 19, 2030	DP			
	9993467*PED	Nov 19, 2030				
<u>POMALIDOMIDE - POMALYST</u>						
N 204026	004 10555939	May 19, 2030	DP		ODE-296	May 14, 2027
	10555939*PED	Nov 19, 2030			ODE-297	May 14, 2027
	8198262	Jun 17, 2025	U-1360		PED	Nov 14, 2027
	8198262	Jun 17, 2025	U-2254		PED	Nov 14, 2027
	8198262*PED	Dec 17, 2025				
	8828427	Jun 21, 2031	DS DP			
	8828427*PED	Dec 21, 2031				
	9993467	May 19, 2030	DP			
	9993467*PED	Nov 19, 2030				

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>PONATINIB HYDROCHLORIDE - ICLUSIG</u>						
N 203469 001	11192895	Dec 12, 2033	U-1700		I-934	Mar 19, 2027
	11192895	Dec 12, 2033	U-1701		ODE-472	Mar 19, 2031
	11192895	Dec 12, 2033	U-1948			
	11192897	Dec 12, 2033	DS U-1700			
	11192897	Dec 12, 2033	DS U-1701			
	11192897	Dec 12, 2033	DS U-1948			
	11384086	Dec 12, 2033	DS DP U-1700			
	11384086	Dec 12, 2033	DS DP U-1701			
	11384086	Dec 12, 2033	DS DP U-1948			
	8114874	Jan 24, 2027	DS DP			
	9029533	Dec 22, 2026	U-1283			
	9029533	Dec 22, 2026	U-1699			
	9029533	Dec 22, 2026	U-1700			
	9029533	Dec 22, 2026	U-1701			
	9029533	Dec 22, 2026	U-836			
	9493470	Dec 12, 2033	DS DP U-1700			
	9493470	Dec 12, 2033	DS DP U-1948			
<u>PONATINIB HYDROCHLORIDE - ICLUSIG</u>						
N 203469 002	11192895	Dec 12, 2033	U-1700		I-934	Mar 19, 2027
	11192895	Dec 12, 2033	U-1701		ODE-472	Mar 19, 2031
	11192895	Dec 12, 2033	U-1948			
	11192897	Dec 12, 2033	DS U-1700			
	11192897	Dec 12, 2033	DS U-1701			
	11192897	Dec 12, 2033	DS U-1948			
	11384086	Dec 12, 2033	DS DP U-1700			
	11384086	Dec 12, 2033	DS DP U-1701			
	11384086	Dec 12, 2033	DS DP U-1948			
	8114874	Jan 24, 2027	DS DP			
	9029533	Dec 22, 2026	U-1283			
	9029533	Dec 22, 2026	U-1699			
	9029533	Dec 22, 2026	U-1700			
	9029533	Dec 22, 2026	U-1701			
	9029533	Dec 22, 2026	U-836			
	9493470	Dec 12, 2033	DS DP U-1700			
	9493470	Dec 12, 2033	DS DP U-1948			
<u>PONATINIB HYDROCHLORIDE - ICLUSIG</u>						
N 203469 003	11192895	Dec 12, 2033	U-1700		I-934	Mar 19, 2027
	11192895	Dec 12, 2033	U-1701		ODE-472	Mar 19, 2031
	11192895	Dec 12, 2033	U-1948			
	11192897	Dec 12, 2033	DS U-1700			
	11192897	Dec 12, 2033	DS U-1701			
	11192897	Dec 12, 2033	DS U-1948			
	11384086	Dec 12, 2033	DS DP U-1700			
	11384086	Dec 12, 2033	DS DP U-1701			
	11384086	Dec 12, 2033	DS DP U-1948			
	8114874	Jan 24, 2027	DS DP			
	9029533	Dec 22, 2026	U-1283			
	9029533	Dec 22, 2026	U-1699			
	9029533	Dec 22, 2026	U-1700			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>PONATINIB HYDROCHLORIDE - ICLUSIG</u>						
N 203469	003	9029533	Dec 22, 2026	U-1701		
		9029533	Dec 22, 2026	U-836		
		9493470	Dec 12, 2033	DS DP U-1700		
		9493470	Dec 12, 2033	DS DP U-1948		
<u>PONATINIB HYDROCHLORIDE - ICLUSIG</u>						
N 203469	004	11192895	Dec 12, 2033	U-1700	I-934	Mar 19, 2027
		11192895	Dec 12, 2033	U-1701	ODE-472	Mar 19, 2031
		11192895	Dec 12, 2033	U-1948		
		11192897	Dec 12, 2033	DS U-1700		
		11192897	Dec 12, 2033	DS U-1701		
		11192897	Dec 12, 2033	DS U-1948		
		11384086	Dec 12, 2033	DS DP U-1700		
		11384086	Dec 12, 2033	DS DP U-1701		
		11384086	Dec 12, 2033	DS DP U-1948		
		8114874	Jan 24, 2027	DS DP		
		9029533	Dec 22, 2026	U-1283		
		9029533	Dec 22, 2026	U-1699		
		9029533	Dec 22, 2026	U-1700		
		9029533	Dec 22, 2026	U-1701		
		9029533	Dec 22, 2026	U-836		
		9493470	Dec 12, 2033	DS DP U-1700		
		9493470	Dec 12, 2033	DS DP U-1948		
<u>PONESIMOD - PONVORY</u>						
N 213498	001	10220023	Dec 10, 2035	U-3103	NCE	Mar 18, 2026
		11951097	Oct 10, 2042	U-3891		
		8273779	Dec 17, 2025	U-2774		
		9062014	May 06, 2032	DS DP U-2774		
		RE43728	Nov 16, 2025	DS DP		
<u>PONESIMOD - PONVORY</u>						
N 213498	002	10220023	Dec 10, 2035	U-3103	NCE	Mar 18, 2026
		11951097	Oct 10, 2042	U-3891		
		8273779	Dec 17, 2025	U-2774		
		9062014	May 06, 2032	DS DP U-2774		
		RE43728	Nov 16, 2025	DS DP		
<u>PONESIMOD - PONVORY</u>						
N 213498	003	10220023	Dec 10, 2035	U-3103	NCE	Mar 18, 2026
		11951097	Oct 10, 2042	U-3891		
		8273779	Dec 17, 2025	U-2774		
		9062014	May 06, 2032	DS DP U-2774		
		RE43728	Nov 16, 2025	DS DP		
<u>PONESIMOD - PONVORY</u>						
N 213498	004	10220023	Dec 10, 2035	U-3103	NCE	Mar 18, 2026
		11951097	Oct 10, 2042	U-3891		
		8273779	Dec 17, 2025	U-2774		
		9062014	May 06, 2032	DS DP U-2774		
		RE43728	Nov 16, 2025	DS DP		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>PONESIMOD - PONVORY</u>						
N 213498 005	10220023	Dec 10, 2035	U-3103		NCE	Mar 18, 2026
	11951097	Oct 10, 2042	U-3891			
	8273779	Dec 17, 2025	U-2774			
	9062014	May 06, 2032	DS DP U-2774			
	RE43728	Nov 16, 2025	DS DP			
<u>PONESIMOD - PONVORY</u>						
N 213498 006	10220023	Dec 10, 2035	U-3103		NCE	Mar 18, 2026
	11951097	Oct 10, 2042	U-3891			
	8273779	Dec 17, 2025	U-2774			
	9062014	May 06, 2032	DS DP U-2774			
	RE43728	Nov 16, 2025	DS DP			
<u>PONESIMOD - PONVORY</u>						
N 213498 007	10220023	Dec 10, 2035	U-3103		NCE	Mar 18, 2026
	11951097	Oct 10, 2042	U-3891			
	8273779	Dec 17, 2025	U-2774			
	9062014	May 06, 2032	DS DP U-2774			
	RE43728	Nov 16, 2025	DS DP			
<u>PONESIMOD - PONVORY</u>						
N 213498 008	10220023	Dec 10, 2035	U-3103		NCE	Mar 18, 2026
	11951097	Oct 10, 2042	U-3891			
	8273779	Dec 17, 2025	U-2774			
	9062014	May 06, 2032	DS DP U-2774			
	RE43728	Nov 16, 2025	DS DP			
<u>PONESIMOD - PONVORY</u>						
N 213498 009	10220023	Dec 10, 2035	U-3103		NCE	Mar 18, 2026
	11951097	Oct 10, 2042	U-3891			
	8273779	Dec 17, 2025	U-2774			
	9062014	May 06, 2032	DS DP U-2774			
	RE43728	Nov 16, 2025	DS DP			
<u>PONESIMOD - PONVORY</u>						
N 213498 010	10220023	Dec 10, 2035	U-3103		NCE	Mar 18, 2026
	11951097	Oct 10, 2042	U-3891			
	8273779	Dec 17, 2025	U-2774			
	9062014	May 06, 2032	DS DP U-2774			
	RE43728	Nov 16, 2025	DS DP			
<u>POSACONAZOLE - NOXAFIL</u>						
N 205053 001					ODE-355	Jun 17, 2028
<u>POSACONAZOLE - NOXAFIL</u>						
N 205596 001	10117951	Mar 13, 2029	DP		ODE-355	Jun 17, 2028
	8410077	Mar 13, 2029	DP			
	9023790	Jul 04, 2031	DP U-1698			
	9023790	Jul 04, 2031	DP U-3160			
	9023790	Jul 04, 2031	DP U-3171			
	9358297	Jun 24, 2031	DP U-3160			
	9358297	Jun 24, 2031	DP U-3171			
	9493582	Feb 27, 2033	DP			
	9750822	Mar 13, 2029	DP			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>POSACONAZOLE - NOXAFIL</u>						
N 205596 001	10117951	Mar 13, 2029	DP		ODE-355	Jun 17, 2028
	8410077	Mar 13, 2029	DP			
	9023790	Jul 04, 2031	DP U-1698			
	9023790	Jul 04, 2031	DP U-3160			
	9023790	Jul 04, 2031	DP U-3171			
	9358297	Jun 24, 2031	DP U-3160			
	9358297	Jun 24, 2031	DP U-3171			
	9493582	Feb 27, 2033	DP			
	9750822	Mar 13, 2029	DP			
<u>POTASSIUM PHOSPHATE, DIBASIC; POTASSIUM PHOSPHATE, MONOBASIC - POTASSIUM PHOSPHATES</u>						
N 212121 001	10632150	Apr 19, 2039	DP U-2789			
<u>PRALATREXATE - FOLOTYN</u>						
N 022468 001	7622470	May 31, 2025	U-1015			
	8299078	May 31, 2025	U-1004			
<u>PRALATREXATE - FOLOTYN</u>						
N 022468 002	7622470	May 31, 2025	U-1015			
	8299078	May 31, 2025	U-1004			
<u>PRALSETINIB - GAVRETO</u>						
N 213721 001	10030005	Nov 01, 2036	DS DP U-2952		NCE	Sep 04, 2025
	10030005	Nov 01, 2036	DS DP U-3002		ODE-318	Sep 04, 2027
	11273160	Apr 03, 2039	U-2952		ODE-340	Dec 01, 2027
	11872192	Apr 03, 2039	U-2952		ODE-341	Dec 01, 2027
	11963958	Apr 03, 2039	U-2828			
<u>PRAMIPEXOLE DIHYDROCHLORIDE - MIRAPEX ER</u>						
N 022421 001	7695734	Apr 26, 2028	DP			
	8679533	Sep 08, 2029	DP U-219			
<u>PRAMIPEXOLE DIHYDROCHLORIDE - MIRAPEX ER</u>						
N 022421 002	7695734	Apr 26, 2028	DP			
	8679533	Sep 08, 2029	DP U-219			
<u>PRAMIPEXOLE DIHYDROCHLORIDE - MIRAPEX ER</u>						
N 022421 003	7695734	Apr 26, 2028	DP			
	8679533	Sep 08, 2029	DP U-219			
<u>PRAMIPEXOLE DIHYDROCHLORIDE - MIRAPEX ER</u>						
N 022421 004	7695734	Apr 26, 2028	DP			
	8679533	Sep 08, 2029	DP U-219			
<u>PRAMIPEXOLE DIHYDROCHLORIDE - MIRAPEX ER</u>						
N 022421 005	7695734	Apr 26, 2028	DP			
	8679533	Sep 08, 2029	DP U-219			
<u>PRAMIPEXOLE DIHYDROCHLORIDE - MIRAPEX ER</u>						
N 022421 006	7695734	Apr 26, 2028	DP			
	8679533	Sep 08, 2029	DP U-219			
<u>PRAMIPEXOLE DIHYDROCHLORIDE - MIRAPEX ER</u>						
N 022421 007	7695734	Apr 26, 2028	DP			
	8679533	Sep 08, 2029	DP U-219			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>PRASTERONE - INTRAROSA</u>						
N 208470	001 8268806	Mar 19, 2031	DP			
	8629129	Aug 07, 2028	DP			
	8957054	Jan 08, 2030	U-1922			
<u>PREDNISOLONE ACETATE - PREDNISOLONE ACETATE</u>						
A 216935	001				CGT	Apr 09, 2025
<u>PREDNISOLONE ACETATE - FLO-PRED</u>						
N 022067	001 7799331	Oct 11, 2028	DP U-1068			
	7799331	Oct 11, 2028	DP U-139			
<u>PREDNISOLONE ACETATE - FLO-PRED</u>						
N 022067	002 7799331	Oct 11, 2028	DP U-1068			
	7799331	Oct 11, 2028	DP U-139			
<u>PREDNISONE - RAYOS</u>						
N 202020	001 9504699	Aug 03, 2027	U-1362			
<u>PREDNISONE - RAYOS</u>						
N 202020	002 9504699	Aug 03, 2027	U-1362			
<u>PREDNISONE - RAYOS</u>						
N 202020	003 8168218	Jan 07, 2028	DP U-1269			
	9504699	Aug 03, 2027	U-1362			
<u>PREGABALIN - LYRICA CR</u>						
N 209501	001 10022447	Nov 02, 2026	U-2136			
	10022447	Nov 02, 2026	U-2137			
	10022447*PED	May 02, 2027				
	8945620	Nov 02, 2026	DP U-2136			
	8945620	Nov 02, 2026	DP U-2137			
	8945620*PED	May 02, 2027				
	9144559	Nov 02, 2026	DP			
	9144559*PED	May 02, 2027				
<u>PREGABALIN - LYRICA CR</u>						
N 209501	002 10022447	Nov 02, 2026	U-2136			
	10022447	Nov 02, 2026	U-2137			
	10022447*PED	May 02, 2027				
	8945620	Nov 02, 2026	DP U-2136			
	8945620	Nov 02, 2026	DP U-2137			
	8945620*PED	May 02, 2027				
	9144559	Nov 02, 2026	DP			
	9144559*PED	May 02, 2027				
<u>PREGABALIN - LYRICA CR</u>						
N 209501	003 10022447	Nov 02, 2026	U-2136			
	10022447	Nov 02, 2026	U-2137			
	10022447*PED	May 02, 2027				
	8945620	Nov 02, 2026	DP U-2136			
	8945620	Nov 02, 2026	DP U-2137			
	8945620*PED	May 02, 2027				
	9144559	Nov 02, 2026	DP			
	9144559*PED	May 02, 2027				

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>PRETOMANID - PRETOMANID</u>						
N 212862	001				NCE	Aug 14, 2024
					ODE-253	Aug 14, 2026
					GAIN	Aug 14, 2029
<u>PROBENECID; SULOPENEM ETZADROXIL - ORLYNVAH</u>						
N 213972	001	11478428	Dec 23, 2039	DP U-4026	NCE	Oct 25, 2029
		11554112	Apr 01, 2039	U-4026	GAIN	Oct 25, 2034
		12109197	Apr 01, 2039	U-4026		
		7795243	Jun 03, 2029	DS DP U-4026		
<u>PROGESTERONE - MILPROSA</u>						
N 201110	001	10537584	Feb 03, 2029	DP		
		10548904	Feb 03, 2029	U-2810		
		8580293	Jan 21, 2030	U-2810		
<u>PROPOFOL - DIPRIVAN</u>						
N 019627	002	8476010*PED	Jun 01, 2025			
<u>PROPRANOLOL HYDROCHLORIDE - HEMANGEOL</u>						
N 205410	001	8338489	Oct 16, 2028	U-1496		
		8987262	Oct 16, 2028	U-1988		
<u>PRUCALOPRIDE SUCCINATE - PRUCALOPRIDE SUCCINATE</u>						
A 218492	001				CGT	Jun 24, 2025
<u>PRUCALOPRIDE SUCCINATE - PRUCALOPRIDE SUCCINATE</u>						
A 218492	002				CGT	Jun 24, 2025
<u>PYRIDOSTIGMINE BROMIDE - PYRIDOSTIGMINE BROMIDE</u>						
N 217604	001	10881617	Jun 18, 2038	DP		
		10925833	Jun 18, 2038	DP		
		10987311	Jun 18, 2038	DP		
		11229606	Jun 18, 2038	DP		
		11478425	Jun 18, 2038	DP U-4012		
		11666536	Jun 18, 2038	DP		
		11911515	Jun 18, 2038	DP		
		12042559	Jun 18, 2038	DP		
<u>QUAZEPAM - DORAL</u>						
N 018708	001	7608616	Jun 03, 2028	U-1012		
<u>QUAZEPAM - DORAL</u>						
N 018708	003	7608616	Jun 03, 2028	U-1012		
<u>QUIZARTINIB DIHYDROCHLORIDE - VANFLYTA</u>						
N 216993	001	7820657	Sep 26, 2028	DS	NCE	Jul 20, 2028
		7968543	Aug 15, 2029	DP U-3661	ODE-437	Jul 20, 2030
		8129374	Mar 16, 2027	U-3661		
		8357690	Feb 26, 2031	U-3661		
		8557810	Mar 16, 2027	DP		
		8836218	Mar 23, 2030	U-3661		
		8865710	Aug 15, 2029	DP		
		8883783	Mar 16, 2027	DS		
		9555040	May 14, 2030	U-3661		
		9585892	Mar 16, 2027	U-3661		
		9675549	Sep 30, 2033	DP		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>QUIZARTINIB DIHYDROCHLORIDE - VANFLYTA</u>						
N 216993 001	7820657	Sep 26, 2028	DS		NCE	Jul 20, 2028
	7968543	Aug 15, 2029	DP U-3661		ODE-437	Jul 20, 2030
	8129374	Mar 16, 2027	U-3661			
	8357690	Feb 26, 2031	U-3661			
	8557810	Mar 16, 2027	DP			
	8836218	Mar 23, 2030	U-3661			
	8865710	Aug 15, 2029	DP			
	8883783	Mar 16, 2027	DS			
	9555040	May 14, 2030	U-3661			
	9585892	Mar 16, 2027	U-3661			
	9675549	Sep 30, 2033	DP			
<u>QUIZARTINIB DIHYDROCHLORIDE - VANFLYTA</u>						
N 216993 002	7820657	Sep 26, 2028	DS		NCE	Jul 20, 2028
	7968543	Aug 15, 2029	DP U-3661		ODE-437	Jul 20, 2030
	8129374	Mar 16, 2027	U-3661			
	8357690	Feb 26, 2031	U-3661			
	8557810	Mar 16, 2027	DP			
	8836218	Mar 23, 2030	U-3661			
	8865710	Aug 15, 2029	DP			
	8883783	Mar 16, 2027	DS			
	9555040	May 14, 2030	U-3661			
	9585892	Mar 16, 2027	U-3661			
	9675549	Sep 30, 2033	DP			
<u>RALTEGRAVIR POTASSIUM - ISENTRESS</u>						
N 022145 001	7754731	Mar 11, 2029	DS DP U-257			
	7754731*PED	Sep 11, 2029				
	8771733	Jun 02, 2030	DS DP U-257			
	8852632	Jan 28, 2028	DS DP U-257			
<u>RALTEGRAVIR POTASSIUM - ISENTRESS HD</u>						
N 022145 002	10772888	Mar 30, 2032			U-1663	
	7754731	Mar 11, 2029	DS DP U-257			
	7754731*PED	Sep 11, 2029				
	8771733	Jun 02, 2030	DS DP U-257			
	9649311	Oct 21, 2030	DP			
	9649311*PED	Apr 21, 2031				
<u>RALTEGRAVIR POTASSIUM - ISENTRESS</u>						
N 203045 001	7754731	Mar 11, 2029	DS DP U-257			
	7754731*PED	Sep 11, 2029				
<u>RALTEGRAVIR POTASSIUM - ISENTRESS</u>						
N 203045 002	7754731	Mar 11, 2029	DS DP U-257			
	7754731*PED	Sep 11, 2029				
	8771733	Jun 02, 2030	DS DP U-257			
<u>RALTEGRAVIR POTASSIUM - ISENTRESS</u>						
N 205786 001	7754731	Mar 11, 2029	DS DP U-257			
	7754731*PED	Sep 11, 2029				

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>RANOLAZINE - ASPRUZYO SPRINKLE</u>						
N 216018	001 11510878	Jan 24, 2038	DP			
<u>RANOLAZINE - ASPRUZYO SPRINKLE</u>						
N 216018	002 10898444	Jan 24, 2038	DP			
	11510878	Jan 24, 2038	DP			
	12161761	Jan 24, 2038	DP			
<u>RASAGILINE MESYLATE - AZILECT</u>						
N 021641	001 7572834	Dec 05, 2026	DP			
	7815942	Aug 27, 2027	DS DP U-219			
<u>RASAGILINE MESYLATE - AZILECT</u>						
N 021641	002 7572834	Dec 05, 2026	DP			
	7815942	Aug 27, 2027	DS DP U-219			
<u>REGADENOSON - LEXISCAN</u>						
N 022161	001 8106183	Feb 02, 2027	DS			
	RE47301	Feb 02, 2027	DP			
<u>REGORAFENIB - STIVARGA</u>						
N 203085	001 8637553	Feb 16, 2031	DS DP			
	8680124	Jun 02, 2030	U-1506			
	9458107	Apr 08, 2031	DP			
	9957232	Jul 09, 2032	DS			
<u>RELUGOLIX - ORGOVYX</u>						
N 214621	001 10350170	Feb 25, 2036	DP		NCE	Dec 18, 2025
	10449191	Sep 29, 2037	U-3020			
	10786501	Sep 29, 2037	U-3020			
	11583526	Sep 29, 2037	U-3020			
	11795178	Sep 27, 2033	DS DP			
	12097198	Sep 29, 2037	U-3020			
	12144809	Sep 29, 2037	U-4035			
	7300935	Jan 28, 2026	DS			
	8058280	Jan 28, 2026	DS DP			
<u>REMDESIVIR - VEKLURY</u>						
N 214787	001 10065958	Sep 16, 2031	DS		D-183	Jan 21, 2025
	10065958*PED	Mar 16, 2032			M-301	Jul 13, 2026
	10675296	Jul 10, 2038	DP		NCE	Oct 22, 2025
	10675296*PED	Jan 10, 2039			NPP	Apr 25, 2025
	10695361	Sep 16, 2036	U-3831		PED	Jul 21, 2025
	10695361	Sep 16, 2036	U-3832		PED	Oct 25, 2025
	10695361*PED	Mar 16, 2037			PED	Apr 22, 2026
	11007208	Sep 16, 2036	U-3831		PED	Jan 13, 2027
	11007208	Sep 16, 2036	U-3832			
	11007208*PED	Mar 16, 2037				
	11266681	Jul 10, 2038	U-3831			
	11266681	Jul 10, 2038	U-3832			
	11266681*PED	Jan 10, 2039				
	11382926	Sep 16, 2036	U-3831			
	11382926	Sep 16, 2036	U-3832			
	11382926*PED	Mar 16, 2037				
	11491169	May 28, 2041	U-3835			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>REMEDESIVIR - VEKLURY</u>						
N 214787 001	11491169	May 28, 2041	U-3836			
	11491169*PED	Nov 28, 2041				
	11492353	Dec 08, 2031	DS			
	11492353*PED	Jun 08, 2032				
	11903953	May 28, 2041	U-3835			
	11903953	May 28, 2041	U-3836			
	11903953*PED	Nov 28, 2041				
	11975012	May 28, 2041	U-3835			
	11975012	May 28, 2041	U-3836			
	11975012*PED	Nov 28, 2041				
	11975017	Jul 10, 2038	DP			
	11975017*PED	Jan 10, 2039				
	8008264	Sep 06, 2029	DS DP			
	8008264*PED	Mar 06, 2030				
	8318682	Apr 22, 2029	DS DP			
	8318682*PED	Oct 22, 2029				
	9724360	Oct 29, 2035	DS DP			
	9724360*PED	Apr 29, 2036				
	9949994	Oct 29, 2035	DS			
	9949994*PED	Apr 29, 2036				
	RE46762	Apr 22, 2029	DS DP			
	RE46762*PED	Oct 22, 2029				
<u>REMEDESIVIR - VEKLURY</u>						
N 214787 002	10065958	Sep 16, 2031	DS		D-183	Jan 21, 2025
	10065958*PED	Mar 16, 2032			M-301	Jul 13, 2026
	10695361	Sep 16, 2036	U-3829		NCE	Oct 22, 2025
	10695361	Sep 16, 2036	U-3830		NPP	Apr 25, 2025
	10695361*PED	Mar 16, 2037			PED	Jul 21, 2025
	11007208	Sep 16, 2036	U-3829		PED	Oct 25, 2025
	11007208	Sep 16, 2036	U-3830		PED	Apr 22, 2026
	11007208*PED	Mar 16, 2037			PED	Jan 13, 2027
	11382926	Sep 16, 2036	U-3829			
	11382926	Sep 16, 2036	U-3830			
	11382926*PED	Mar 16, 2037				
	11491169	May 28, 2041	U-3833			
	11491169	May 28, 2041	U-3834			
	11491169*PED	Nov 28, 2041				
	11492353	Dec 08, 2031	DS			
	11492353*PED	Jun 08, 2032				
	11903953	May 28, 2041	U-3833			
	11903953	May 28, 2041	U-3834			
	11903953*PED	Nov 28, 2041				
	11975012	May 28, 2041	U-3833			
	11975012	May 28, 2041	U-3834			
	11975012*PED	Nov 28, 2041				
	8008264	Sep 06, 2029	DS DP			
	8008264*PED	Mar 06, 2030				
	8318682	Apr 22, 2029	DS DP			
	8318682*PED	Oct 22, 2029				

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>REMEDESIVIR - VEKLURY</u>						
N 214787 002	9724360	Oct 29, 2035	DS DP			
	9724360*PED	Apr 29, 2036				
	9949994	Oct 29, 2035	DS			
	9949994*PED	Apr 29, 2036				
	RE46762	Apr 22, 2029	DS DP			
	RE46762*PED	Oct 22, 2029				
<u>REMIMAZOLAM BESYLATE - BYFAVO</u>						
N 212295 001	10052334	Nov 07, 2031	U-2968		NCE	Oct 06, 2025
	10195210	Nov 07, 2031	U-2968			
	10342800	Nov 07, 2031	U-2968			
	10472365	Jul 10, 2027	U-2968			
	10722522	Nov 07, 2031	U-2968			
	10961250	Jul 10, 2027	DP U-2968			
	9561236	Apr 30, 2033	U-2968			
	9737547	Nov 07, 2031	U-2968			
	9777007	Jul 10, 2027	DP			
	9827251	Jan 13, 2034	U-2968			
	9914738	Jul 10, 2027	DP			
<u>REPOTRECTINIB - AUGTYRO</u>						
N 218213 001	10294242	Jul 05, 2036	DS		NCE	Nov 15, 2028
	11452725	Jul 24, 2036	U-3755		ODE-455	Nov 15, 2030
	11452725	Jul 24, 2036	U-3961		ODE-483	Jun 13, 2031
	9714258	Jan 23, 2035	DS DP		ODE-488	Jun 13, 2031
<u>REPOTRECTINIB - AUGTYRO</u>						
N 218213 002	10294242	Jul 05, 2036	DS		NCE	Nov 15, 2028
	11452725	Jul 24, 2036	U-3755		ODE*	Nov 15, 2030
	11452725	Jul 24, 2036	U-3961		ODE-488	Jun 13, 2031
	9714258	Jan 23, 2035	DS DP			
<u>RESMETIROM - REZDIFFRA</u>						
N 217785 001	10376517	Sep 17, 2033	U-3861		NCE	Mar 14, 2029
	11564926	Sep 17, 2033	DS DP U-3861			
	11986481	Sep 17, 2033	U-3861			
	7452882	Sep 12, 2026	DS DP			
	9266861	Sep 17, 2033	DS DP			
<u>RESMETIROM - REZDIFFRA</u>						
N 217785 002	10376517	Sep 17, 2033	U-3861		NCE	Mar 14, 2029
	11564926	Sep 17, 2033	DS DP U-3861			
	11986481	Sep 17, 2033	U-3861			
	7452882	Sep 12, 2026	DS DP			
	9266861	Sep 17, 2033	DS DP			
<u>RESMETIROM - REZDIFFRA</u>						
N 217785 003	10376517	Sep 17, 2033	U-3861		NCE	Mar 14, 2029
	11564926	Sep 17, 2033	DS DP U-3861			
	11986481	Sep 17, 2033	U-3861			
	7452882	Sep 12, 2026	DS DP			
	9266861	Sep 17, 2033	DS DP			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>RETAPAMULIN - ALTABAX</u>						
N 022055	001 7875630	Feb 14, 2027	DS			
<u>REVEFENACIN - YUPELRI</u>						
N 210598	001 10106503	Mar 10, 2025		U-2440		
	10343995	Mar 10, 2025		U-2440		
	10550081	Jul 14, 2030	DS			
	11008289	Jul 14, 2030		U-2440		
	11247969	Mar 10, 2025	DP			
	11484531	Oct 23, 2039		U-2440		
	11691948	Jul 14, 2030	DP			
	11858898	Jul 14, 2030	DS DP	U-2440		
	12048692	Aug 29, 2039		U-2440		
	7288657	Oct 31, 2028	DS			
	7491736	Mar 10, 2025		U-2440		
	7521041	Mar 10, 2025		U-2440		
	7550595	Mar 10, 2025	DP			
	7585879	Mar 10, 2025	DS DP	U-2440		
	7910608	Mar 10, 2025	DS DP			
	8034946	Mar 10, 2025	DP			
	8053448	Mar 10, 2025		U-2440		
	8273894	Mar 10, 2025	DP			
	8541451	Aug 25, 2031	DS			
	9765028	Jul 14, 2030	DS			
<u>REVUMENIB CITRATE - REVUFORJ</u>						
N 218944	001 10683302	Jun 08, 2037	DS DP		NCE	Nov 15, 2029
	11479557	Jun 08, 2037	DP U-4045		ODE-502	Nov 15, 2031
<u>REVUMENIB CITRATE - REVUFORJ</u>						
N 218944	002 10683302	Jun 08, 2037	DS DP		NCE	Nov 15, 2029
	11479557	Jun 08, 2037	DP U-4045		ODE-502	Nov 15, 2031
<u>REVUMENIB CITRATE - REVUFORJ</u>						
N 218944	003 10683302	Jun 08, 2037	DS DP		NCE	Nov 15, 2029
	11479557	Jun 08, 2037	DP U-4045		ODE-502	Nov 15, 2031
<u>REZAFUNGIN ACETATE - REZZAYO</u>						
N 217417	001 10702573	Mar 14, 2033	DS DP U-3566		NCE	Mar 22, 2028
	11197909	Jul 14, 2038	DS DP U-3566		ODE-426	Mar 22, 2030
	11654196	Mar 02, 2032	DS DP U-3566		GAIN	Mar 22, 2033
	11712459	Mar 15, 2037	DP U-3566		GAIN	Mar 22, 2035
	11819533	Jul 11, 2038	DS DP U-3566			
	8722619	Mar 02, 2032	DS DP U-3566			
	9526835	Mar 14, 2033	DS DP U-3566			
<u>RIBOCICLIB SUCCINATE - KISOALI</u>						
N 209092	001 10799506	Apr 14, 2036	DP		I-950	Sep 17, 2027
	12064434	Apr 14, 2036	DP			
	8324225	Jun 17, 2028	DS DP			
	8415355	Mar 13, 2031	DS DP			
	8685980	May 25, 2030	DS DP			
	8962630	Dec 09, 2029		U-1981		
	8962630	Dec 09, 2029		U-2355		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>RIBOCICLIB SUCCINATE - KISQALI</u>						
N 209092 001	8962630	Dec 09, 2029	U-2356			
	8962630	Dec 09, 2029	U-3265			
	8962630	Dec 09, 2029	U-3266			
	8962630	Dec 09, 2029	U-3975			
	8962630	Dec 09, 2029	U-3999			
	9193732	Nov 09, 2031	DS DP			
	9416136	Aug 20, 2029	U-1981			
	9416136	Aug 20, 2029	U-2355			
	9416136	Aug 20, 2029	U-2356			
	9416136	Aug 20, 2029	U-3265			
	9416136	Aug 20, 2029	U-3266			
	9416136	Aug 20, 2029	U-3975			
	9416136	Aug 20, 2029	U-3999			
	9868739	Nov 09, 2031	U-1981			
	9868739	Nov 09, 2031	U-2355			
	9868739	Nov 09, 2031	U-2356			
	9868739	Nov 09, 2031	U-3265			
	9868739	Nov 09, 2031	U-3266			
	9868739	Nov 09, 2031	U-3975			
	9868739	Nov 09, 2031	U-3999			
<u>RIFAMYCIN SODIUM - AEMCOLO</u>						
N 210910 001	8263120	May 03, 2025	DP		NCE	Nov 16, 2023
	8486446	May 03, 2025	DP		GAIN	Nov 16, 2028
	8529945	May 03, 2025	DP			
	8741948	May 03, 2025	DP U-2448			
<u>RIFAXIMIN - XIFAXAN</u>						
N 021361 001	10703763	Feb 27, 2026	U-1708			
	10703763	Feb 27, 2026	U-2847			
	10703763	Feb 27, 2026	U-2848			
	7906542	Jun 01, 2025	DS DP			
	7928115	Jul 24, 2029	U-1121			
	8193196	Sep 02, 2027	DS DP			
	8518949	Feb 27, 2026	DP			
	8741904	Feb 27, 2026	DS U-1526			
	9271968	Feb 27, 2026	DP			
<u>RIFAXIMIN - XIFAXAN</u>						
N 021361 002	10314828	Jul 24, 2029	U-1481			
	10335397	Jul 24, 2029	U-2579			
	10456384	Feb 26, 2029	U-2643			
	10456384	Feb 26, 2029	U-2644			
	10703763	Feb 27, 2026	U-1708			
	10703763	Feb 27, 2026	U-2847			
	10703763	Feb 27, 2026	U-2848			
	10709694	Jul 24, 2029	U-2579			
	10765667	Feb 26, 2029	U-2643			
	10765667	Feb 26, 2029	U-2644			
	11564912	Feb 26, 2029	U-3511			
	11564912	Feb 26, 2029	U-3512			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>RIFAXIMIN - XIFAXAN</u>						
N 021361	002 11779571	Feb 26, 2029	U-3706			
	7906542	Jun 01, 2025	DS DP			
	7915275	Feb 23, 2025	U-1707			
	7915275	Feb 23, 2025	U-1708			
	8193196	Sep 02, 2027	DS DP U-1707			
	8193196	Sep 02, 2027	DS DP U-1708			
	8309569	Jul 18, 2029	U-1707			
	8309569	Jul 18, 2029	U-1708			
	8518949	Feb 27, 2026	DP			
	8642573	Oct 02, 2029	U-1481			
	8741904	Feb 27, 2026	DS U-1526			
	8741904	Feb 27, 2026	DS U-1707			
	8741904	Feb 27, 2026	DS U-1708			
	8829017	Jul 24, 2029	U-1562			
	8946252	Jul 24, 2029	U-1481			
	8969398	Oct 02, 2029	U-1481			
	9271968	Feb 27, 2026	DP			
	9421195	Jul 24, 2029	U-1481			
	9629828	Jul 24, 2029	U-1994			
<u>RILPIVIRINE HYDROCHLORIDE - EDURANT</u>						
N 202022	001 7125879	Apr 21, 2025	DS DP U-1153			
	7125879	Apr 21, 2025	DS DP U-1307			
	7125879	Apr 21, 2025	DS DP U-1740			
	7125879	Apr 21, 2025	DS DP U-3353			
	7125879	Apr 21, 2025	DS DP U-3874			
	7125879*PED	Oct 21, 2025				
<u>RILPIVIRINE HYDROCHLORIDE - EDURANT PED</u>						
N 219016	001 11065198	Oct 23, 2037	DP			
	7125879	Apr 21, 2025	DS DP U-3874			
<u>RILUZOLE - TIGLUTIK KIT</u>						
N 209080	001 8765150	Mar 12, 2029	DP U-2401			
<u>RIMEGEPANT SULFATE - NURTEC ODT</u>						
N 212728	001 11083724	Mar 25, 2039	DP U-2718		NCE	Feb 27, 2025
	11083724	Mar 25, 2039	DP U-3142			
	8314117	Mar 09, 2030	DS DP U-2718			
	8314117	Mar 09, 2030	DS DP U-3142			
	8759372	Feb 25, 2033	DS DP			
<u>RIOCIGUAT - ADEMPAS</u>						
N 204819	001 10662188	Feb 18, 2034	DS DP U-2834			
	10662188	Feb 18, 2034	DS DP U-2835			
	11203593	Feb 18, 2034	DS DP U-2834			
	11203593	Feb 18, 2034	DS DP U-2835			
	7173037	Dec 04, 2026	DS DP			
<u>RIOCIGUAT - ADEMPAS</u>						
N 204819	002 10662188	Feb 18, 2034	DS DP U-2834			
	10662188	Feb 18, 2034	DS DP U-2835			
	11203593	Feb 18, 2034	DS DP U-2834			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>RIOCIGUAT - ADEMPAS</u>						
N 204819 002	11203593	Feb 18, 2034	DS DP U-2835			
	7173037	Dec 04, 2026	DS DP			
<u>RIOCIGUAT - ADEMPAS</u>						
N 204819 003	10662188	Feb 18, 2034	DS DP U-2834			
	10662188	Feb 18, 2034	DS DP U-2835			
	11203593	Feb 18, 2034	DS DP U-2834			
	11203593	Feb 18, 2034	DS DP U-2835			
	7173037	Dec 04, 2026	DS DP			
<u>RIOCIGUAT - ADEMPAS</u>						
N 204819 004	10662188	Feb 18, 2034	DS DP U-2834			
	10662188	Feb 18, 2034	DS DP U-2835			
	11203593	Feb 18, 2034	DS DP U-2834			
	11203593	Feb 18, 2034	DS DP U-2835			
	7173037	Dec 04, 2026	DS DP			
<u>RIOCIGUAT - ADEMPAS</u>						
N 204819 005	10662188	Feb 18, 2034	DS DP U-2834			
	10662188	Feb 18, 2034	DS DP U-2835			
	11203593	Feb 18, 2034	DS DP U-2834			
	11203593	Feb 18, 2034	DS DP U-2835			
	7173037	Dec 04, 2026	DS DP			
<u>RIPRETINIB - QINLOCK</u>						
N 213973 001	10966966	Aug 12, 2040	U-3153		NCE	May 15, 2025
	11185535	Dec 30, 2040	DP		ODE-298	May 15, 2027
	11266635	Aug 12, 2040	U-3330			
	11344536	Aug 12, 2040	U-3381			
	11395818	Dec 30, 2040	DP			
	11426390	Aug 12, 2040	U-3416			
	11433056	Aug 12, 2040	U-3423			
	11529336	Aug 12, 2040	U-3382			
	11534432	Aug 12, 2040	U-3442			
	11576903	Dec 30, 2040	DP			
	11576904	Aug 12, 2040	U-3537			
	11612591	Dec 30, 2040	DP			
	11779572	Oct 06, 2042	U-3714			
	11793795	Dec 30, 2040	DP			
	11801237	Dec 30, 2040	DP U-3219			
	11813251	Aug 12, 2040	U-3750			
	11844788	Dec 30, 2040	DP			
	11850240	Dec 30, 2040	DP			
	11850241	Dec 30, 2040	DP			
	11896585	Dec 30, 2040	DP			
	11903933	Dec 30, 2040	DP			
	11911370	Dec 30, 2040	DP			
	11918564	Dec 30, 2040	DP			
	11969414	Feb 08, 2041	U-3897			
	11969415	Dec 30, 2040	U-3219			
	12023325	Aug 12, 2040	U-3960			
	12023326	Aug 12, 2040	U-3959			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>RIPRETINIB - QINLOCK</u>						
N 213973	001	12023327	Aug 12, 2040	U-3958		
		12023328	Dec 30, 2040	U-3957		
		12059410	Aug 12, 2040	U-3153		
		12059411	Aug 12, 2040	U-3971		
		12064422	Dec 30, 2040	U-3219		
		8188113	Jul 27, 2030	DS DP		
		8461179	May 15, 2034	DS DP		
		RE48731	Jun 07, 2032	U-3219		
<u>RISDIPLAM - EVRYSDI</u>						
N 213535	001	11534444	Oct 04, 2038	U-1943	M-270	Oct 03, 2026
		11827646	Jan 25, 2036	U-1943	NCE	Aug 07, 2025
		11938136	Nov 08, 2036	DP	NPP	May 27, 2025
		12122789	Apr 15, 2041	DS DP	ODE-334	Aug 07, 2027
		9586955	Feb 08, 2033	DS DP	ODE-400	May 27, 2029
		9969754	May 11, 2035	DS DP U-1943		
<u>RISEDRONATE SODIUM - ATELVIA</u>						
N 022560	001	7645459	Jan 09, 2028	DP U-662		
		7645460	Jan 09, 2028	DP U-662		
		8246989	Jan 16, 2026	DP		
<u>RISPERIDONE - PERSERIS KIT</u>						
N 210655	001	10010612	Feb 13, 2028	DP		
		10058554	Sep 26, 2026	U-2363		
		10376590	Feb 13, 2028	U-2608		
		10406160	Jun 26, 2026	DP U-2608		
		11013809	Feb 13, 2028	DP U-3135		
		11110093	Nov 05, 2026	DP U-3135		
		11712475	Feb 13, 2028	U-3135		
		9180197	Feb 13, 2028	DP		
		9186413	Feb 13, 2028	U-543		
		9597402	Sep 26, 2026	DP		
<u>RISPERIDONE - PERSERIS KIT</u>						
N 210655	002	10010612	Feb 13, 2028	DP		
		10058554	Sep 26, 2026	U-2363		
		10376590	Feb 13, 2028	U-2608		
		10406160	Jun 26, 2026	DP U-2608		
		11013809	Feb 13, 2028	DP U-3135		
		11110093	Nov 05, 2026	DP U-3135		
		11712475	Feb 13, 2028	U-3135		
		9180197	Feb 13, 2028	DP		
		9186413	Feb 13, 2028	U-543		
		9597402	Sep 26, 2026	DP		
<u>RISPERIDONE - RYKINDO</u>						
N 212849	001	10098882	Apr 10, 2032	DP U-3513		
		10406161	Apr 10, 2032	DP U-3513		
		11110094	Apr 10, 2032	DP		
		9446135	Apr 10, 2032	DP U-3513		
		9532991	Apr 10, 2032	DP U-3513		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>RISPERIDONE - RYKINDO</u>						
N 212849	002	10098882	Apr 10, 2032	DP U-3513		
		10406161	Apr 10, 2032	DP U-3513		
		11110094	Apr 10, 2032	DP		
		9446135	Apr 10, 2032	DP U-3513		
		9532991	Apr 10, 2032	DP U-3513		
<u>RISPERIDONE - RYKINDO</u>						
N 212849	003	10098882	Apr 10, 2032	DP U-3513		
		10406161	Apr 10, 2032	DP U-3513		
		11110094	Apr 10, 2032	DP		
		9446135	Apr 10, 2032	DP U-3513		
		9532991	Apr 10, 2032	DP U-3513		
<u>RISPERIDONE - RYKINDO</u>						
N 212849	004	10098882	Apr 10, 2032	DP U-3513		
		10406161	Apr 10, 2032	DP U-3513		
		11110094	Apr 10, 2032	DP		
		9446135	Apr 10, 2032	DP U-3513		
		9532991	Apr 10, 2032	DP U-3513		
<u>RISPERIDONE - UZEDY</u>						
N 213586	001	10736965	Jan 12, 2025	DP	NP	Apr 28, 2026
		12128132	Sep 11, 2040	DP U-543		
		8221778	Nov 12, 2027	DP U-543		
		8741327	Nov 12, 2027	DP U-543		
		8802127	Jan 12, 2025	DP		
		9023897	Apr 05, 2033	DP		
		9439905	Jan 12, 2025	DP U-543		
		9717799	Jan 12, 2025	DP		
		9895447	Jan 12, 2025	DP		
		9925268	Jan 12, 2025	DP		
<u>RISPERIDONE - UZEDY</u>						
N 213586	002	10736965	Jan 12, 2025	DP	NP	Apr 28, 2026
		12128132	Sep 11, 2040	DP U-543		
		8221778	Nov 12, 2027	DP U-543		
		8741327	Nov 12, 2027	DP U-543		
		8802127	Jan 12, 2025	DP		
		9023897	Apr 05, 2033	DP		
		9439905	Jan 12, 2025	DP U-543		
		9717799	Jan 12, 2025	DP		
		9895447	Jan 12, 2025	DP		
		9925268	Jan 12, 2025	DP		
<u>RISPERIDONE - UZEDY</u>						
N 213586	003	10736965	Jan 12, 2025	DP	NP	Apr 28, 2026
		12128132	Sep 11, 2040	DP U-543		
		8221778	Nov 12, 2027	DP U-543		
		8741327	Nov 12, 2027	DP U-543		
		8802127	Jan 12, 2025	DP		
		9023897	Apr 05, 2033	DP		
		9439905	Jan 12, 2025	DP U-543		
		9717799	Jan 12, 2025	DP		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>RISPERIDONE - UZEDY</u>						
N 213586	003 9895447	Jan 12, 2025	DP			
	9925268	Jan 12, 2025	DP			
<u>RISPERIDONE - UZEDY</u>						
N 213586	004 10736965	Jan 12, 2025	DP		NP	Apr 28, 2026
	12128132	Sep 11, 2040	DP U-543			
	8221778	Nov 12, 2027	DP U-543			
	8741327	Nov 12, 2027	DP U-543			
	8802127	Jan 12, 2025	DP			
	9023897	Apr 05, 2033	DP			
	9439905	Jan 12, 2025	DP U-543			
	9717799	Jan 12, 2025	DP			
	9895447	Jan 12, 2025	DP			
	9925268	Jan 12, 2025	DP			
<u>RISPERIDONE - UZEDY</u>						
N 213586	005 10736965	Jan 12, 2025	DP		NP	Apr 28, 2026
	12128132	Sep 11, 2040	DP U-543			
	8221778	Nov 12, 2027	DP U-543			
	8741327	Nov 12, 2027	DP U-543			
	8802127	Jan 12, 2025	DP			
	9023897	Apr 05, 2033	DP			
	9439905	Jan 12, 2025	DP U-543			
	9717799	Jan 12, 2025	DP			
	9895447	Jan 12, 2025	DP			
	9925268	Jan 12, 2025	DP			
<u>RISPERIDONE - UZEDY</u>						
N 213586	006 10736965	Jan 12, 2025	DP		NP	Apr 28, 2026
	12128132	Sep 11, 2040	DP U-543			
	8221778	Nov 12, 2027	DP U-543			
	8741327	Nov 12, 2027	DP U-543			
	8802127	Jan 12, 2025	DP			
	9023897	Apr 05, 2033	DP			
	9439905	Jan 12, 2025	DP U-543			
	9717799	Jan 12, 2025	DP			
	9895447	Jan 12, 2025	DP			
	9925268	Jan 12, 2025	DP			
<u>RISPERIDONE - UZEDY</u>						
N 213586	007 10736965	Jan 12, 2025	DP		NP	Apr 28, 2026
	12128132	Sep 11, 2040	DP U-543			
	8221778	Nov 12, 2027	DP U-543			
	8741327	Nov 12, 2027	DP U-543			
	8802127	Jan 12, 2025	DP			
	9023897	Apr 05, 2033	DP			
	9439905	Jan 12, 2025	DP U-543			
	9717799	Jan 12, 2025	DP			
	9895447	Jan 12, 2025	DP			
	9925268	Jan 12, 2025	DP			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>RISPERIDONE - RISVAN</u>						
N 214835	001	10058504	May 31, 2031	DP	NP	Mar 29, 2027
		10085936	May 31, 2031	DP		
		10182982	May 31, 2031	U-3943		
		10195138	May 31, 2031	U-3943		
		10335366	May 31, 2031	DP		
		10463607	May 31, 2031	DP		
		10881605	May 31, 2031	DP		
		11007139	May 31, 2031	U-3942		
		11173110	May 31, 2031	U-3941		
		11241377	May 31, 2031	U-3939		
		11752092	May 31, 2031	U-3941		
		11752093	May 31, 2031	DP		
		11752094	May 31, 2031	U-3942		
		11759416	May 31, 2031	U-3939		
<u>RISPERIDONE - RISVAN</u>						
N 214835	002	10058504	May 31, 2031	DP	NP	Mar 29, 2027
		10085936	May 31, 2031	DP		
		10182982	May 31, 2031	U-3943		
		10195138	May 31, 2031	U-3943		
		10335366	May 31, 2031	DP		
		10463607	May 31, 2031	DP		
		10881605	May 31, 2031	DP		
		11007139	May 31, 2031	U-3942		
		11173110	May 31, 2031	U-3941		
		11241377	May 31, 2031	U-3939		
		11752092	May 31, 2031	U-3941		
		11752093	May 31, 2031	DP		
		11752094	May 31, 2031	U-3942		
		11759416	May 31, 2031	U-3939		
<u>RITLECITINIB TOSYLATE - LITFULO</u>						
N 215830	001	12077533	Dec 03, 2034	U-3994	NCE	Jun 23, 2028
		12116368	Oct 17, 2041	DS		
		9617258	Dec 03, 2034	DS DP		
<u>RIVAROXABAN - XARELTO</u>						
N 022406	001	7157456*PED	Feb 28, 2025		I-867	Aug 23, 2024
		9415053*PED	May 13, 2025		PED	Feb 23, 2025
		9539218	Feb 17, 2034	U-1957		
		9539218	Feb 17, 2034	U-2143		
		9539218	Feb 17, 2034	U-2641		
		9539218	Feb 17, 2034	U-3288		
		9539218*PED	Aug 17, 2034			
<u>RIVAROXABAN - XARELTO</u>						
N 022406	002	7157456*PED	Feb 28, 2025		I-867	Aug 23, 2024
		9415053*PED	May 13, 2025		PED	Feb 23, 2025
		9539218	Feb 17, 2034	U-1953		
		9539218	Feb 17, 2034	U-3289		
		9539218*PED	Aug 17, 2034			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>RIVAROXABAN - XARELTO</u>						
N 022406 003	7157456*PED	Feb 28, 2025			I-867	Aug 23, 2024
	9415053*PED	May 13, 2025			PED	Feb 23, 2025
	9539218	Feb 17, 2034	U-1953			
	9539218	Feb 17, 2034	U-1954			
	9539218	Feb 17, 2034	U-1955			
	9539218	Feb 17, 2034	U-3285			
	9539218*PED	Aug 17, 2034				
<u>RIVAROXABAN - XARELTO</u>						
N 022406 004	10828310	Jan 31, 2039	U-3207		I-867	Aug 23, 2024
	10828310	Jan 31, 2039	U-3208		PED	Feb 23, 2025
	10828310*PED	Jul 31, 2039				
	7157456*PED	Feb 28, 2025				
	9415053*PED	May 13, 2025				
<u>RIVAROXABAN - XARELTO</u>						
N 215859 001	7157456*PED	Feb 28, 2025			NP	Dec 20, 2024
					PED	Jun 20, 2025
<u>RIZATRIPTAN BENZOATE - RIZAFILM</u>						
N 205394 001	9301948	Jul 30, 2034	DS DP			
<u>ROFLUMILAST - ZORYVE</u>						
N 215985 001	10940142	Jun 07, 2037	DP		NP	Jul 29, 2025
	11129818	Aug 25, 2037	U-3408		NPP	Oct 05, 2026
	11129818	Aug 25, 2037	U-3712			
	11793796	Jun 07, 2037	DP			
	11819496	Jun 07, 2037	U-3748			
	11992480	Jun 07, 2037	U-3748			
	12005051	Jun 07, 2037	U-3748			
	12005052	Jun 07, 2037	DP			
	12011437	Jun 07, 2037	DP			
	12016848	Jun 07, 2037	DP			
	12042487	Jun 07, 2037	DP			
	9884050	Jun 07, 2037	DP			
	9907788	Jun 07, 2037	U-3408			
	9907788	Jun 07, 2037	U-3712			
<u>ROFLUMILAST - ZORYVE</u>						
N 215985 002	10940142	Jun 07, 2037	DP		NS	Jul 09, 2027
	11129818	Aug 25, 2037	U-3970			
	11793796	Jun 07, 2037	DP			
	11819496	Jun 07, 2037	U-3970			
	11992480	Jun 07, 2037	U-3970			
	12005051	Jun 07, 2037	U-3970			
	12005052	Jun 07, 2037	DP			
	12011437	Jun 07, 2037	DP			
	12016848	Jun 07, 2037	DP			
	12042487	Jun 07, 2037	DP			
	9884050	Jun 07, 2037	DP			
	9907788	Jun 07, 2037	U-3970			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>ROFLUMILAST - ZORYVE</u>						
N 217242	001	10940142	Jun 07, 2037	DP	NP	Dec 15, 2026
		11129818	Aug 25, 2037	U-3773		
		11707454	Dec 03, 2041	U-3773		
		11793796	Jun 07, 2037	DP		
		11819496	Jun 07, 2037	U-3773		
		11992480	Jun 07, 2037	U-3773		
		12005051	Jun 07, 2037	U-3773		
		12005052	Jun 07, 2037	DP		
		12011437	Jun 07, 2037	DP		
		12016848	Jun 07, 2037	DP		
		12042487	Jun 07, 2037	DP		
		9884050	Jun 07, 2037	DP		
		9907788	Jun 07, 2037	U-3773		
<u>ROLAPITANT HYDROCHLORIDE - VARUBI</u>						
N 206500	001	7049320	Aug 19, 2028	DS DP	U-1741	
		7563801	Apr 04, 2027	DP		
		7981905	Apr 04, 2027	U-1741		
		8178550	Apr 04, 2027	DS DP		
		8361500	Oct 09, 2029	DP		
		8404702	Apr 04, 2027	U-1741		
		8470842	Jan 18, 2029	U-1741		
<u>ROLAPITANT HYDROCHLORIDE - VARUBI</u>						
N 208399	001	7049320	Aug 19, 2028	DS DP	U-1741	
		7981905	Apr 04, 2027	U-1741		
		8178550	Apr 04, 2027	DS DP		
		8404702	Apr 04, 2027	U-1741		
		8470842	Jan 18, 2029	U-1741		
		9101615	Jul 14, 2032	U-1741		
<u>ROPIVACAINE HYDROCHLORIDE - NAROPIN</u>						
N 020533	006	7828787	Oct 18, 2025	DP		
		7857802	Nov 28, 2026	DP		
<u>ROPIVACAINE HYDROCHLORIDE - NAROPIN</u>						
N 020533	007	7828787	Oct 18, 2025	DP		
		7857802	Nov 28, 2026	DP		
<u>ROSUVASTATIN CALCIUM - EZALLOR SPRINKLE</u>						
N 208647	001	10413543	Feb 12, 2036	DP		
<u>ROSUVASTATIN CALCIUM - EZALLOR SPRINKLE</u>						
N 208647	002	10413543	Feb 12, 2036	DP		
<u>ROSUVASTATIN CALCIUM - EZALLOR SPRINKLE</u>						
N 208647	003	10413543	Feb 12, 2036	DP		
<u>ROSUVASTATIN CALCIUM - EZALLOR SPRINKLE</u>						
N 208647	004	10413543	Feb 12, 2036	DP		
<u>ROTIGOTINE - NEUPRO</u>						
N 021829	001	10130589	Dec 22, 2030	DP		
		10350174	Dec 22, 2030	DP		
		8246979	Sep 01, 2027	DP	U-1272	

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>ROTIGOTINE - NEUPRO</u>						
N 021829 001	8246979	Sep 01, 2027	DP U-1273			
	8246980	Nov 27, 2025	DP			
	9925150	Mar 01, 2032	DP			
<u>ROTIGOTINE - NEUPRO</u>						
N 021829 002	10130589	Dec 22, 2030	DP			
	10350174	Dec 22, 2030	DP			
	8246979	Sep 01, 2027	DP U-1272			
	8246979	Sep 01, 2027	DP U-1273			
	8246980	Nov 27, 2025	DP			
	9925150	Mar 01, 2032	DP			
<u>ROTIGOTINE - NEUPRO</u>						
N 021829 003	10130589	Dec 22, 2030	DP			
	10350174	Dec 22, 2030	DP			
	8246979	Sep 01, 2027	DP U-1272			
	8246979	Sep 01, 2027	DP U-1273			
	8246980	Nov 27, 2025	DP			
	9925150	Mar 01, 2032	DP			
<u>ROTIGOTINE - NEUPRO</u>						
N 021829 004	10130589	Dec 22, 2030	DP			
	10350174	Dec 22, 2030	DP			
	8246979	Sep 01, 2027	DP U-1272			
	8246979	Sep 01, 2027	DP U-1273			
	8246980	Nov 27, 2025	DP			
	9925150	Mar 01, 2032	DP			
<u>ROTIGOTINE - NEUPRO</u>						
N 021829 005	10130589	Dec 22, 2030	DP			
	10350174	Dec 22, 2030	DP			
	8246979	Sep 01, 2027	DP U-1272			
	8246979	Sep 01, 2027	DP U-1273			
	8246980	Nov 27, 2025	DP			
	9925150	Mar 01, 2032	DP			
<u>ROTIGOTINE - NEUPRO</u>						
N 021829 006	10130589	Dec 22, 2030	DP			
	10350174	Dec 22, 2030	DP			
	8246979	Sep 01, 2027	DP U-1272			
	8246979	Sep 01, 2027	DP U-1273			
	8246980	Nov 27, 2025	DP			
	9925150	Mar 01, 2032	DP			
<u>RUCAPARIB CAMSYLATE - RUBRACA</u>						
N 209115 001	10130636	Aug 17, 2035	U-2012		ODE-168	Apr 06, 2025
	10130636	Aug 17, 2035	U-2101			
	10130636	Aug 17, 2035	U-2273			
	10130636	Aug 17, 2035	U-2830			
	10278974	Feb 10, 2031	DP			
	8071579	Aug 12, 2027	U-2012			
	8071579	Aug 12, 2027	U-2273			
	8071579	Aug 12, 2027	U-2830			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>RUCAPARIB CAMSYLATE - RUBRACA</u>						
N 209115 001	8143241	Aug 12, 2027	U-2012			
	8143241	Aug 12, 2027	U-2273			
	8143241	Aug 12, 2027	U-2830			
	8754072	Feb 10, 2031	DS DP			
	8859562	Aug 04, 2031	U-2012			
	8859562	Aug 04, 2031	U-2273			
	8859562	Aug 04, 2031	U-2830			
	9045487	Feb 10, 2031	DS DP			
	9861638	Feb 10, 2031	U-2012			
	9861638	Feb 10, 2031	U-2273			
	9987285	Aug 17, 2035	DP			
<u>RUCAPARIB CAMSYLATE - RUBRACA</u>						
N 209115 002	10130636	Aug 17, 2035	U-2012		ODE-168	Apr 06, 2025
	10130636	Aug 17, 2035	U-2101			
	10130636	Aug 17, 2035	U-2273			
	10130636	Aug 17, 2035	U-2830			
	10278974	Feb 10, 2031	DP			
	8071579	Aug 12, 2027	U-2012			
	8071579	Aug 12, 2027	U-2273			
	8071579	Aug 12, 2027	U-2830			
	8143241	Aug 12, 2027	U-2012			
	8143241	Aug 12, 2027	U-2273			
	8143241	Aug 12, 2027	U-2830			
	8754072	Feb 10, 2031	DS DP			
	8859562	Aug 04, 2031	U-2012			
	8859562	Aug 04, 2031	U-2273			
	8859562	Aug 04, 2031	U-2830			
	9045487	Feb 10, 2031	DS DP			
	9861638	Feb 10, 2031	U-2012			
	9861638	Feb 10, 2031	U-2273			
	9987285	Aug 17, 2035	DP			
<u>RUCAPARIB CAMSYLATE - RUBRACA</u>						
N 209115 003	10130636	Aug 17, 2035	U-2012		ODE-168	Apr 06, 2025
	10130636	Aug 17, 2035	U-2101			
	10130636	Aug 17, 2035	U-2273			
	10130636	Aug 17, 2035	U-2830			
	10278974	Feb 10, 2031	DP			
	8071579	Aug 12, 2027	U-2012			
	8071579	Aug 12, 2027	U-2273			
	8071579	Aug 12, 2027	U-2830			
	8143241	Aug 12, 2027	U-2012			
	8143241	Aug 12, 2027	U-2273			
	8143241	Aug 12, 2027	U-2830			
	8754072	Feb 10, 2031	DS DP			
	8859562	Aug 04, 2031	U-2012			
	8859562	Aug 04, 2031	U-2273			
	8859562	Aug 04, 2031	U-2830			
	9045487	Feb 10, 2031	DS DP			
	9861638	Feb 10, 2031	U-2012			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>RUCAPARIB CAMSYLATE - RUBRACA</u>						
N 209115 003	9861638	Feb 10, 2031	U-2273			
	9987285	Aug 17, 2035	DP			
<u>RUXOLITINIB PHOSPHATE - JAKAFI</u>						
N 202192 001	10016429	Jun 12, 2028	U-3226		I-872	Sep 22, 2024
	10016429	Jun 12, 2028	U-3230		M-285	Dec 19, 2025
	10016429*PED	Dec 12, 2028			ODE-238	May 24, 2026
	7598257	Dec 24, 2027	DS DP U-3227		ODE-373	Sep 22, 2028
	7598257	Dec 24, 2027	DS DP U-3228		PED	Mar 22, 2025
	7598257*PED	Jun 24, 2028			PED	Jun 19, 2026
	8415362	Dec 24, 2027	DS DP		PED	Nov 24, 2026
	8415362*PED	Jun 24, 2028			PED	Mar 22, 2029
	8722693	Jun 12, 2028	DS DP			
	8722693*PED	Dec 12, 2028				
	8822481	Jun 12, 2028	U-1573			
	8822481	Jun 12, 2028	U-3226			
	8822481	Jun 12, 2028	U-3227			
	8822481	Jun 12, 2028	U-3228			
	8822481	Jun 12, 2028	U-3230			
	8822481*PED	Dec 12, 2028				
	8829013	Jun 12, 2028	U-1201			
	8829013	Jun 12, 2028	U-1622			
	8829013	Jun 12, 2028	U-3227			
	8829013	Jun 12, 2028	U-3228			
	8829013*PED	Dec 12, 2028				
	9079912	Dec 12, 2026	U-3226			
	9079912	Dec 12, 2026	U-3227			
	9079912	Dec 12, 2026	U-3228			
	9079912	Dec 12, 2026	U-3230			
	9079912*PED	Jun 12, 2027				
	9814722	Dec 12, 2026	U-3226			
	9814722	Dec 12, 2026	U-3230			
	9814722*PED	Jun 12, 2027				
<u>RUXOLITINIB PHOSPHATE - JAKAFI</u>						
N 202192 002	10016429	Jun 12, 2028	U-3226		I-872	Sep 22, 2024
	10016429	Jun 12, 2028	U-3230		M-285	Dec 19, 2025
	10016429*PED	Dec 12, 2028			ODE-238	May 24, 2026
	7598257	Dec 24, 2027	DS DP U-3227		ODE-373	Sep 22, 2028
	7598257	Dec 24, 2027	DS DP U-3228		PED	Mar 22, 2025
	7598257*PED	Jun 24, 2028			PED	Jun 19, 2026
	8415362	Dec 24, 2027	DS DP		PED	Nov 24, 2026
	8415362*PED	Jun 24, 2028			PED	Mar 22, 2029
	8722693	Jun 12, 2028	DS DP			
	8722693*PED	Dec 12, 2028				
	8822481	Jun 12, 2028	U-1573			
	8822481	Jun 12, 2028	U-3226			
	8822481	Jun 12, 2028	U-3227			
	8822481	Jun 12, 2028	U-3228			
	8822481	Jun 12, 2028	U-3230			
	8822481*PED	Dec 12, 2028				

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>RUXOLITINIB PHOSPHATE - JAKAFI</u>						
N 202192 002	8829013	Jun 12, 2028	U-1201			
	8829013	Jun 12, 2028	U-1622			
	8829013	Jun 12, 2028	U-3227			
	8829013	Jun 12, 2028	U-3228			
	8829013*PED	Dec 12, 2028				
	9079912	Dec 12, 2026	U-3226			
	9079912	Dec 12, 2026	U-3227			
	9079912	Dec 12, 2026	U-3228			
	9079912	Dec 12, 2026	U-3230			
	9079912*PED	Jun 12, 2027				
	9814722	Dec 12, 2026	U-3226			
	9814722	Dec 12, 2026	U-3230			
	9814722*PED	Jun 12, 2027				
<u>RUXOLITINIB PHOSPHATE - JAKAFI</u>						
N 202192 003	10016429	Jun 12, 2028	U-3226		M-285	Dec 19, 2025
	10016429	Jun 12, 2028	U-3230		ODE-238	May 24, 2026
	10016429*PED	Dec 12, 2028			ODE-373	Sep 22, 2028
	7598257	Dec 24, 2027	DS DP U-3227		PED	Jun 19, 2026
	7598257	Dec 24, 2027	DS DP U-3228		PED	Nov 24, 2026
	7598257*PED	Jun 24, 2028			PED	Mar 22, 2029
	8415362	Dec 24, 2027	DS DP			
	8415362*PED	Jun 24, 2028				
	8722693	Jun 12, 2028	DS DP			
	8722693*PED	Dec 12, 2028				
	8822481	Jun 12, 2028	U-1573			
	8822481	Jun 12, 2028	U-3226			
	8822481	Jun 12, 2028	U-3227			
	8822481	Jun 12, 2028	U-3228			
	8822481	Jun 12, 2028	U-3230			
	8822481*PED	Dec 12, 2028				
	8829013	Jun 12, 2028	U-1201			
	8829013	Jun 12, 2028	U-1622			
	8829013	Jun 12, 2028	U-3227			
	8829013	Jun 12, 2028	U-3228			
	8829013*PED	Dec 12, 2028				
	9079912	Dec 12, 2026	U-3226			
	9079912	Dec 12, 2026	U-3227			
	9079912	Dec 12, 2026	U-3228			
	9079912	Dec 12, 2026	U-3230			
	9079912*PED	Jun 12, 2027				
	9814722	Dec 12, 2026	U-3226			
	9814722	Dec 12, 2026	U-3230			
	9814722*PED	Jun 12, 2027				
<u>RUXOLITINIB PHOSPHATE - JAKAFI</u>						
N 202192 004	10016429	Jun 12, 2028	U-3226		M-285	Dec 19, 2025
	10016429	Jun 12, 2028	U-3230		ODE-238	May 24, 2026
	10016429*PED	Dec 12, 2028			ODE-373	Sep 22, 2028
	7598257	Dec 24, 2027	DS DP U-3227		PED	Jun 19, 2026
	7598257	Dec 24, 2027	DS DP U-3228		PED	Nov 24, 2026

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>RUXOLITINIB PHOSPHATE - JAKAFI</u>						
N 202192 004	7598257*PED	Jun 24, 2028			PED	Mar 22, 2029
	8415362	Dec 24, 2027	DS DP			
	8415362*PED	Jun 24, 2028				
	8722693	Jun 12, 2028	DS DP			
	8722693*PED	Dec 12, 2028				
	8822481	Jun 12, 2028		U-1573		
	8822481	Jun 12, 2028		U-3226		
	8822481	Jun 12, 2028		U-3227		
	8822481	Jun 12, 2028		U-3228		
	8822481	Jun 12, 2028		U-3230		
	8822481*PED	Dec 12, 2028				
	8829013	Jun 12, 2028		U-1201		
	8829013	Jun 12, 2028		U-1622		
	8829013	Jun 12, 2028		U-3227		
	8829013	Jun 12, 2028		U-3228		
	8829013*PED	Dec 12, 2028				
	9079912	Dec 12, 2026		U-3226		
	9079912	Dec 12, 2026		U-3227		
	9079912	Dec 12, 2026		U-3228		
	9079912	Dec 12, 2026		U-3230		
	9079912*PED	Jun 12, 2027				
	9814722	Dec 12, 2026		U-3226		
	9814722	Dec 12, 2026		U-3230		
	9814722*PED	Jun 12, 2027				
<u>RUXOLITINIB PHOSPHATE - JAKAFI</u>						
N 202192 005	10016429	Jun 12, 2028		U-3226	M-285	Dec 19, 2025
	10016429	Jun 12, 2028		U-3230	ODE-238	May 24, 2026
	10016429*PED	Dec 12, 2028			ODE-373	Sep 22, 2028
	7598257	Dec 24, 2027	DS DP	U-3227	PED	Jun 19, 2026
	7598257	Dec 24, 2027	DS DP	U-3228	PED	Nov 24, 2026
	7598257*PED	Jun 24, 2028			PED	Mar 22, 2029
	8415362	Dec 24, 2027	DS DP			
	8415362*PED	Jun 24, 2028				
	8722693	Jun 12, 2028	DS DP			
	8722693*PED	Dec 12, 2028				
	8822481	Jun 12, 2028		U-1573		
	8822481	Jun 12, 2028		U-3226		
	8822481	Jun 12, 2028		U-3227		
	8822481	Jun 12, 2028		U-3228		
	8822481	Jun 12, 2028		U-3230		
	8822481*PED	Dec 12, 2028				
	8829013	Jun 12, 2028		U-1201		
	8829013	Jun 12, 2028		U-1622		
	8829013	Jun 12, 2028		U-3227		
	8829013	Jun 12, 2028		U-3228		
	8829013*PED	Dec 12, 2028				
	9079912	Dec 12, 2026		U-3226		
	9079912	Dec 12, 2026		U-3227		
	9079912	Dec 12, 2026		U-3228		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>RUXOLITINIB PHOSPHATE - JAKAFI</u>						
N 202192 005	9079912	Dec 12, 2026	U-3230			
	9079912*PED	Jun 12, 2027				
	9814722	Dec 12, 2026	U-3226			
	9814722	Dec 12, 2026	U-3230			
	9814722*PED	Jun 12, 2027				
<u>RUXOLITINIB PHOSPHATE - OPZELURA</u>						
N 215309 001	10610530	Jun 12, 2028	U-3229		I-896	Jul 18, 2025
	10610530	Jun 12, 2028	U-3404		NP	Sep 21, 2024
	10610530*PED	Dec 12, 2028			PED	Mar 21, 2025
	10639310	Dec 12, 2026	U-3229		PED	Jan 18, 2026
	10639310*PED	Jun 12, 2027				
	10758543	May 20, 2031	DP			
	10758543*PED	Nov 20, 2031				
	10869870	May 20, 2031	U-3229			
	10869870	May 20, 2031	U-3404			
	10869870*PED	Nov 20, 2031				
	11219624	May 20, 2031	DP U-3229			
	11219624*PED	Nov 20, 2031				
	11510923	Sep 04, 2040	U-3505			
	11571425	May 20, 2031	DP			
	11590136	May 20, 2031	U-3229			
	11590136	May 20, 2031	U-3404			
	11590137	Sep 04, 2040	U-3505			
	11590138	Jun 10, 2040	U-3551			
	11602536	May 05, 2041	U-3550			
	7598257	Dec 24, 2027	DS DP			
	7598257*PED	Jun 24, 2028				
	8415362	Dec 24, 2027	DS DP			
	8415362*PED	Jun 24, 2028				
	8722693	Jun 12, 2028	DS DP			
	8722693*PED	Dec 12, 2028				
	8822481	Jun 12, 2028	U-3229			
	8822481	Jun 12, 2028	U-3404			
	8822481*PED	Dec 12, 2028				
	9079912	Dec 12, 2026	U-3229			
	9079912	Dec 12, 2026	U-3404			
	9079912*PED	Jun 12, 2027				
	9974790	Dec 12, 2026	U-3229			
	9974790	Dec 12, 2026	U-3404			
	9974790*PED	Jun 12, 2027				
<u>SACUBITRIL; VALSARTAN - ENTRESTO</u>						
N 207620 001	11058667	May 09, 2036	U-3170			
	11135192	Aug 22, 2033	U-3084			
	8101659	Jan 15, 2025	DP			
	8101659*PED	Jul 15, 2025				
	8877938	May 27, 2027	DS DP			
	8877938*PED	Nov 27, 2027				
	9388134	Nov 08, 2026	U-1723			
	9388134*PED	May 08, 2027				

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>SACUBITRIL; VALSARTAN - ENTRESTO</u>						
N 207620	001	9517226	Aug 22, 2033			U-3084
		9937143	Aug 22, 2033			U-3084
<u>SACUBITRIL; VALSARTAN - ENTRESTO</u>						
N 207620	002	11058667	May 09, 2036			U-3170
		11135192	Aug 22, 2033			U-3084
		8101659	Jan 15, 2025	DP		
		8101659*PED	Jul 15, 2025			
		8877938	May 27, 2027	DS DP		
		8877938*PED	Nov 27, 2027			
		9388134	Nov 08, 2026			U-1723
		9388134*PED	May 08, 2027			
		9517226	Aug 22, 2033			U-3084
		9937143	Aug 22, 2033			U-3084
<u>SACUBITRIL; VALSARTAN - ENTRESTO</u>						
N 207620	003	11058667	May 09, 2036			U-3170
		11135192	Aug 22, 2033			U-3084
		8101659	Jan 15, 2025	DP		
		8101659*PED	Jul 15, 2025			
		8877938	May 27, 2027	DS DP		
		8877938*PED	Nov 27, 2027			
		9388134	Nov 08, 2026			U-1723
		9388134*PED	May 08, 2027			
		9517226	Aug 22, 2033			U-3084
		9937143	Aug 22, 2033			U-3084
<u>SACUBITRIL; VALSARTAN - ENTRESTO SPRINKLE</u>						
N 218591	001	10722471	Feb 02, 2037	DP		U-3896
		8101659	Jan 15, 2025	DP		
		8877938	May 27, 2027	DS DP		
		9388134	Nov 08, 2026			U-1723
<u>SACUBITRIL; VALSARTAN - ENTRESTO SPRINKLE</u>						
N 218591	002	10722471	Feb 02, 2037	DP		U-3896
		8101659	Jan 15, 2025	DP		
		8877938	May 27, 2027	DS DP		
		9388134	Nov 08, 2026			U-1723
<u>SAFINAMIDE MESYLATE - XADAGO</u>						
N 207145	001	8076515	Dec 10, 2028	DS DP		U-1993
		8278485	Jun 08, 2027	DS		U-1993
		8283380	Mar 21, 2031			U-1993
<u>SAFINAMIDE MESYLATE - XADAGO</u>						
N 207145	002	8076515	Dec 10, 2028	DS DP		U-1993
		8278485	Jun 08, 2027	DS		U-1993
		8283380	Mar 21, 2031			U-1993
<u>SAPROPTERIN DIHYDROCHLORIDE - KUVAN</u>						
N 022181	001	7566462	Nov 16, 2025	DP		
		7566462*PED	May 16, 2026			
		7566714*PED	May 17, 2025			
		7612073*PED	May 17, 2025			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>SAPROPTERIN DIHYDROCHLORIDE - KUVAN</u>						
N 022181	001	7727987*PED	May 17, 2025			
		8003126	Nov 16, 2025			
		8003126*PED	May 16, 2026			
		8067416*PED	May 17, 2025			
		8318745*PED	May 17, 2025			
		9433624*PED	May 17, 2025			
		RE43797*PED	May 17, 2025			
<u>SAPROPTERIN DIHYDROCHLORIDE - KUVAN</u>						
N 205065	001	7566714*PED	May 17, 2025			
		7612073*PED	May 17, 2025			
		8067416*PED	May 17, 2025			
		9216178	Nov 01, 2032	DP		
		9216178*PED	May 01, 2033			
		9433624*PED	May 17, 2025			
		RE43797*PED	May 17, 2025			
<u>SAPROPTERIN DIHYDROCHLORIDE - KUVAN</u>						
N 205065	002	7566714*PED	May 17, 2025			
		7612073*PED	May 17, 2025			
		8067416*PED	May 17, 2025			
		9216178	Nov 01, 2032	DP		
		9216178*PED	May 01, 2033			
		9433624*PED	May 17, 2025			
		RE43797*PED	May 17, 2025			
<u>SARECYCLINE HYDROCHLORIDE - SEYSARA</u>						
N 209521	001	8318706	May 01, 2031	DS DP	U-2405	
		8513223	Dec 07, 2029		U-2406	
		9255068	Feb 09, 2033	DS DP	U-2407	
		9255068	Feb 09, 2033	DS DP	U-2408	
		9481639	Aug 10, 2028		U-2409	
<u>SARECYCLINE HYDROCHLORIDE - SEYSARA</u>						
N 209521	002	8318706	May 01, 2031	DS DP	U-2405	
		8513223	Dec 07, 2029		U-2406	
		9255068	Feb 09, 2033	DS DP	U-2407	
		9255068	Feb 09, 2033	DS DP	U-2408	
		9481639	Aug 10, 2028		U-2409	
<u>SARECYCLINE HYDROCHLORIDE - SEYSARA</u>						
N 209521	003	8318706	May 01, 2031	DS DP	U-2405	
		8513223	Dec 07, 2029		U-2406	
		9255068	Feb 09, 2033	DS DP	U-2407	
		9255068	Feb 09, 2033	DS DP	U-2408	
		9481639	Aug 10, 2028		U-2409	
<u>SAXAGLIPTIN HYDROCHLORIDE - ONGLYZA</u>						
N 022350	001	7951400	Nov 30, 2028	DP		
<u>SAXAGLIPTIN HYDROCHLORIDE - ONGLYZA</u>						
N 022350	002	7951400	Nov 30, 2028	DP		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>SECNIDAZOLE - SOLOSEC</u>						
N 209363 001	10335390	Sep 04, 2035	DP U-2583		NCE	Sep 15, 2022
	10335390	Sep 04, 2035	DP U-3270		NPP	Jan 26, 2025
	10682338	Sep 04, 2035	DP U-2583		GAIN	Sep 15, 2027
	10682338	Sep 04, 2035	DP U-3270			
	10849884	Sep 04, 2035	DP U-2583			
	10849884	Sep 04, 2035	DP U-3169			
	10849884	Sep 04, 2035	DP U-3270			
	10849884	Sep 04, 2035	DP U-3302			
	10857133	Sep 04, 2035	DP U-2583			
	10857133	Sep 04, 2035	DP U-3270			
	11000507	Sep 04, 2035	DP U-2583			
	11000507	Sep 04, 2035	DP U-3169			
	11000507	Sep 04, 2035	DP U-3270			
	11000507	Sep 04, 2035	DP U-3302			
	11000508	Sep 04, 2035	DP U-2583			
	11000508	Sep 04, 2035	DP U-3169			
	11000508	Sep 04, 2035	DP U-3270			
	11000508	Sep 04, 2035	DP U-3302			
	11020377	Sep 04, 2035	DP U-2583			
	11020377	Sep 04, 2035	DP U-3169			
	11020377	Sep 04, 2035	DP U-3270			
	11020377	Sep 04, 2035	DP U-3302			
	11324721	Sep 04, 2035	DP U-2583			
	11324721	Sep 04, 2035	DP U-3169			
	11324721	Sep 04, 2035	DP U-3270			
	11324721	Sep 04, 2035	DP U-3302			
	11602522	Sep 04, 2035	DP U-3169			
	11602522	Sep 04, 2035	DP U-3302			
	11684607	Sep 16, 2035	DP U-2583			
	11684607	Sep 16, 2035	DP U-3169			
	11684607	Sep 16, 2035	DP U-3270			
	11684607	Sep 16, 2035	DP U-3302			
<u>SELADELPAR LYSINE - LIVDELZI</u>						
N 217899 001	10272058	Mar 19, 2035	U-1854		NCE	Aug 14, 2029
	11406611	Mar 19, 2035	U-1854		ODE-486	Aug 14, 2031
	11596614	Mar 19, 2035	U-1854			
	7301050	Aug 02, 2025	DS DP			
	7709682	Sep 13, 2026	DS			
	9486428	Mar 19, 2035	U-1854			
<u>SELENIOS ACID - SELENIOS ACID</u>						
N 209379 001	11998565	Jul 01, 2041	DP			
	12150957	Jul 01, 2041	U-4039			
<u>SELENIOS ACID - SELENIOS ACID</u>						
N 209379 002	11998565	Jul 01, 2041	DP			
	12150957	Jul 01, 2041	U-4039			
<u>SELENIOS ACID - SELENIOS ACID</u>						
N 209379 003	11998565	Jul 01, 2041	DP			
	12150957	Jul 01, 2041	U-4039			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>SELENIOS ACID - SELENIOS ACID</u>						
N 209379	003 11998565	Jul 01, 2041	DP			
	12150957	Jul 01, 2041	U-4039			
<u>SELEXIPAG - UPTRAVI</u>						
N 207947	001 10821108	Dec 01, 2036	DP U-2992			
	10828298	Dec 01, 2036	DP U-2991			
	7205302	Oct 31, 2026	DS DP U-1797			
	8791122	Aug 01, 2030	DS DP			
	9173881	Aug 12, 2029	U-1798			
	9284280	Jun 25, 2030	U-1831			
<u>SELEXIPAG - UPTRAVI</u>						
N 207947	002 10821108	Dec 01, 2036	DP U-2992			
	10828298	Dec 01, 2036	DP U-2991			
	7205302	Oct 31, 2026	DS DP U-1797			
	8791122	Aug 01, 2030	DS DP			
	9173881	Aug 12, 2029	U-1798			
	9284280	Jun 25, 2030	U-1831			
<u>SELEXIPAG - UPTRAVI</u>						
N 207947	003 10821108	Dec 01, 2036	DP U-2992			
	10828298	Dec 01, 2036	DP U-2991			
	7205302	Oct 31, 2026	DS DP U-1797			
	8791122	Aug 01, 2030	DS DP			
	9173881	Aug 12, 2029	U-1798			
	9284280	Jun 25, 2030	U-1831			
<u>SELEXIPAG - UPTRAVI</u>						
N 207947	004 10821108	Dec 01, 2036	DP U-2992			
	10828298	Dec 01, 2036	DP U-2991			
	7205302	Oct 31, 2026	DS DP U-1797			
	8791122	Aug 01, 2030	DS DP			
	9173881	Aug 12, 2029	U-1798			
	9284280	Jun 25, 2030	U-1831			
<u>SELEXIPAG - UPTRAVI</u>						
N 207947	005 10821108	Dec 01, 2036	DP U-2992			
	10828298	Dec 01, 2036	DP U-2991			
	7205302	Oct 31, 2026	DS DP U-1797			
	8791122	Aug 01, 2030	DS DP			
	9173881	Aug 12, 2029	U-1798			
	9284280	Jun 25, 2030	U-1831			
<u>SELEXIPAG - UPTRAVI</u>						
N 207947	006 10821108	Dec 01, 2036	DP U-2992			
	10828298	Dec 01, 2036	DP U-2991			
	7205302	Oct 31, 2026	DS DP U-1797			
	8791122	Aug 01, 2030	DS DP			
	9173881	Aug 12, 2029	U-1798			
	9284280	Jun 25, 2030	U-1831			
<u>SELEXIPAG - UPTRAVI</u>						
N 207947	007 10821108	Dec 01, 2036	DP U-2992			
	10828298	Dec 01, 2036	DP U-2991			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>SELEXIPAG - UPTRAVI</u>						
N 207947	007 7205302	Oct 31, 2026	DS DP U-1797			
	8791122	Aug 01, 2030	DS DP			
	9173881	Aug 12, 2029		U-1798		
	9284280	Jun 25, 2030		U-1831		
<u>SELEXIPAG - UPTRAVI</u>						
N 207947	008 10821108	Dec 01, 2036		DP U-2992		
	10828298	Dec 01, 2036		DP U-2991		
	7205302	Oct 31, 2026	DS DP U-1797			
	8791122	Aug 01, 2030	DS DP			
	9173881	Aug 12, 2029		U-1798		
	9284280	Jun 25, 2030		U-1831		
<u>SELEXIPAG - UPTRAVI</u>						
N 214275	001 7205302	Oct 31, 2026	DS DP U-1797			
	8791122	Aug 01, 2030	DS DP			
	9173881	Aug 12, 2029		U-1798		
	9284280	Jun 25, 2030		U-1831		
<u>SELINEXOR - XPOVIO</u>						
N 212306	001 10519139	Aug 14, 2035	DS DP U-2584		ODE-257	Jul 03, 2026
	10519139	Aug 14, 2035	DS DP U-2855		ODE-310	Jun 22, 2027
	10519139	Aug 14, 2035	DS DP U-3018		ODE-346	Dec 18, 2027
	10544108	Jul 26, 2032		U-2584		
	10544108	Jul 26, 2032		U-3018		
	11034660	Jul 26, 2032		U-2584		
	11034660	Jul 26, 2032		U-3018		
	11746102	Aug 14, 2035		U-2584		
	11746102	Aug 14, 2035		U-2855		
	11746102	Aug 14, 2035		U-3018		
	11753401	Aug 14, 2035		DP U-2584		
	11753401	Aug 14, 2035		DP U-2855		
	11753401	Aug 14, 2035		DP U-3018		
	11787771	Jul 26, 2032		U-2584		
	11787771	Jul 26, 2032		U-2855		
	11787771	Jul 26, 2032		U-3018		
	11807629	Aug 14, 2035	DS DP			
	8999996	Jul 03, 2033	DS DP			
	9079865	Jul 26, 2032		U-2584		
	9079865	Jul 26, 2032		U-2855		
	9079865	Jul 26, 2032		U-3018		
	9714226	Jul 26, 2032	DS DP			
<u>SELINEXOR - XPOVIO</u>						
N 212306	002 10519139	Aug 14, 2035	DS DP U-2584		ODE*	Jul 03, 2026
	10519139	Aug 14, 2035	DS DP U-2855		ODE*	Jun 22, 2027
	10519139	Aug 14, 2035	DS DP U-3018		ODE*	Dec 18, 2027
	10544108	Jul 26, 2032		U-2584		
	10544108	Jul 26, 2032		U-3018		
	11034660	Jul 26, 2032		U-2584		
	11034660	Jul 26, 2032		U-3018		
	11746102	Aug 14, 2035		U-2584		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>SELINEXOR - XPOVIO</u>						
N 212306 002	11746102	Aug 14, 2035	U-2855			
	11746102	Aug 14, 2035	U-3018			
	11753401	Aug 14, 2035	DP U-2584			
	11753401	Aug 14, 2035	DP U-2855			
	11753401	Aug 14, 2035	DP U-3018			
	11787771	Jul 26, 2032	U-2584			
	11787771	Jul 26, 2032	U-2855			
	11787771	Jul 26, 2032	U-3018			
	11807629	Aug 14, 2035	DS DP			
	8999996	Jul 03, 2033	DS DP			
	9079865	Jul 26, 2032	U-2584			
	9079865	Jul 26, 2032	U-2855			
	9079865	Jul 26, 2032	U-3018			
	9714226	Jul 26, 2032	DS DP			
<u>SELINEXOR - XPOVIO</u>						
N 212306 003	10519139	Aug 14, 2035	DS DP U-2584		ODE*	Jul 03, 2026
	10519139	Aug 14, 2035	DS DP U-2855		ODE*	Jun 22, 2027
	10519139	Aug 14, 2035	DS DP U-3018		ODE*	Dec 18, 2027
	10544108	Jul 26, 2032	U-2584			
	10544108	Jul 26, 2032	U-3018			
	11034660	Jul 26, 2032	U-2584			
	11034660	Jul 26, 2032	U-3018			
	11746102	Aug 14, 2035	U-2584			
	11746102	Aug 14, 2035	U-2855			
	11746102	Aug 14, 2035	U-3018			
	11753401	Aug 14, 2035	DP U-2584			
	11753401	Aug 14, 2035	DP U-2855			
	11753401	Aug 14, 2035	DP U-3018			
	11787771	Jul 26, 2032	U-2584			
	11787771	Jul 26, 2032	U-2855			
	11787771	Jul 26, 2032	U-3018			
	11807629	Aug 14, 2035	DS DP			
	8999996	Jul 03, 2033	DS DP			
	9079865	Jul 26, 2032	U-2584			
	9079865	Jul 26, 2032	U-2855			
	9079865	Jul 26, 2032	U-3018			
	9714226	Jul 26, 2032	DS DP			
<u>SELINEXOR - XPOVIO</u>						
N 212306 004	10519139	Aug 14, 2035	DS DP U-2584		ODE*	Jul 03, 2026
	10519139	Aug 14, 2035	DS DP U-2855		ODE*	Jun 22, 2027
	10519139	Aug 14, 2035	DS DP U-3018		ODE*	Dec 18, 2027
	10544108	Jul 26, 2032	U-2584			
	10544108	Jul 26, 2032	U-3018			
	11034660	Jul 26, 2032	U-2584			
	11034660	Jul 26, 2032	U-3018			
	11746102	Aug 14, 2035	U-2584			
	11746102	Aug 14, 2035	U-2855			
	11746102	Aug 14, 2035	U-3018			
	11753401	Aug 14, 2035	DP U-2584			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>SELINEXOR - XPOVIO</u>						
N 212306 004	11753401	Aug 14, 2035	DP U-2855			
	11753401	Aug 14, 2035	DP U-3018			
	11787771	Jul 26, 2032	U-2584			
	11787771	Jul 26, 2032	U-2855			
	11787771	Jul 26, 2032	U-3018			
	11807629	Aug 14, 2035	DS DP			
	8999996	Jul 03, 2033	DS DP			
	9079865	Jul 26, 2032	U-2584			
	9079865	Jul 26, 2032	U-2855			
	9079865	Jul 26, 2032	U-3018			
	9714226	Jul 26, 2032	DS DP			
<u>SELPERCATINIB - RETEVMO</u>						
N 213246 001	10112942	Oct 10, 2037	DS DP		NCE	May 08, 2025
	10137124	Oct 10, 2037	U-3450		NPP	May 29, 2027
	10137124	Oct 10, 2037	U-3949		ODE-301	May 08, 2027
	10137124	Oct 10, 2037	U-3950		ODE-302	May 08, 2027
	10137124	Oct 10, 2037	U-3951		ODE-303	May 08, 2027
	10172851	Oct 10, 2037	U-3450		ODE-409	Sep 21, 2029
	10172851	Oct 10, 2037	U-3949		ODE-412	Sep 21, 2029
	10172851	Oct 10, 2037	U-3950		ODE-484	May 29, 2031
	10172851	Oct 10, 2037	U-3951		ODE-485	May 29, 2031
	10584124	Oct 10, 2038	DS U-3450		ODE-487	May 29, 2031
	10584124	Oct 10, 2038	DS U-3949			
	10584124	Oct 10, 2038	DS U-3950			
	10584124	Oct 10, 2038	DS U-3951			
	10786489	Oct 10, 2038	DP U-3450			
	10786489	Oct 10, 2038	DP U-3949			
	10786489	Oct 10, 2038	DP U-3950			
	10786489	Oct 10, 2038	DP U-3951			
	12138250	Oct 10, 2038	U-3949			
	12138250	Oct 10, 2038	U-3950			
	12138250	Oct 10, 2038	U-3951			
<u>SELPERCATINIB - RETEVMO</u>						
N 213246 002	10112942	Oct 10, 2037	DS DP		NCE	May 08, 2025
	10137124	Oct 10, 2037	U-3450		NPP	May 29, 2027
	10137124	Oct 10, 2037	U-3949		ODE-301	May 08, 2027
	10137124	Oct 10, 2037	U-3950		ODE-302	May 08, 2027
	10137124	Oct 10, 2037	U-3951		ODE-303	May 08, 2027
	10172851	Oct 10, 2037	U-3450		ODE-409	Sep 21, 2029
	10172851	Oct 10, 2037	U-3949		ODE-412	Sep 21, 2029
	10172851	Oct 10, 2037	U-3950		ODE-484	May 29, 2031
	10172851	Oct 10, 2037	U-3951		ODE-485	May 29, 2031
	10584124	Oct 10, 2038	DS U-3450		ODE-487	May 29, 2031
	10584124	Oct 10, 2038	DS U-3949			
	10584124	Oct 10, 2038	DS U-3950			
	10584124	Oct 10, 2038	DS U-3951			
	10786489	Oct 10, 2038	DP U-3450			
	10786489	Oct 10, 2038	DP U-3949			
	10786489	Oct 10, 2038	DP U-3950			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>SELPERCATINIB - RETEVMO</u>						
N 213246 002	10786489	Oct 10, 2038	DP U-3951			
	12138250	Oct 10, 2038	U-3949			
	12138250	Oct 10, 2038	U-3950			
	12138250	Oct 10, 2038	U-3951			
<u>SELPERCATINIB - RETEVMO</u>						
N 218160 001	10112942	Oct 10, 2037	DS DP		NCE	May 08, 2025
	10137124	Oct 10, 2037	U-3450			
	10137124	Oct 10, 2037	U-3949			
	10137124	Oct 10, 2037	U-3950			
	10137124	Oct 10, 2037	U-3951			
	10172851	Oct 10, 2037	U-3450			
	10172851	Oct 10, 2037	U-3949			
	10172851	Oct 10, 2037	U-3950			
	10172851	Oct 10, 2037	U-3951			
	12138250	Oct 10, 2038	U-3949			
	12138250	Oct 10, 2038	U-3950			
	12138250	Oct 10, 2038	U-3951			
<u>SELPERCATINIB - RETEVMO</u>						
N 218160 002	10112942	Oct 10, 2037	DS DP		NCE	May 08, 2025
	10137124	Oct 10, 2037	U-3450			
	10137124	Oct 10, 2037	U-3949			
	10137124	Oct 10, 2037	U-3950			
	10137124	Oct 10, 2037	U-3951			
	10172851	Oct 10, 2037	U-3450			
	10172851	Oct 10, 2037	U-3949			
	10172851	Oct 10, 2037	U-3950			
	10172851	Oct 10, 2037	U-3951			
	12138250	Oct 10, 2038	U-3949			
	12138250	Oct 10, 2038	U-3950			
	12138250	Oct 10, 2038	U-3951			
<u>SELPERCATINIB - RETEVMO</u>						
N 218160 003	10112942	Oct 10, 2037	DS DP		NCE	May 08, 2025
	10137124	Oct 10, 2037	U-3450			
	10137124	Oct 10, 2037	U-3949			
	10137124	Oct 10, 2037	U-3950			
	10137124	Oct 10, 2037	U-3951			
	10172851	Oct 10, 2037	U-3450			
	10172851	Oct 10, 2037	U-3949			
	10172851	Oct 10, 2037	U-3950			
	10172851	Oct 10, 2037	U-3951			
	12138250	Oct 10, 2038	U-3949			
	12138250	Oct 10, 2038	U-3950			
	12138250	Oct 10, 2038	U-3951			
<u>SELPERCATINIB - RETEVMO</u>						
N 218160 004	10112942	Oct 10, 2037	DS DP		NCE	May 08, 2025
	10137124	Oct 10, 2037	U-3450			
	10137124	Oct 10, 2037	U-3949			
	10137124	Oct 10, 2037	U-3950			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>SELPERCATINIB - RETEVMO</u>						
N 218160 004	10137124	Oct 10, 2037	U-3951			
	10172851	Oct 10, 2037	U-3450			
	10172851	Oct 10, 2037	U-3949			
	10172851	Oct 10, 2037	U-3950			
	10172851	Oct 10, 2037	U-3951			
	12138250	Oct 10, 2038	U-3949			
	12138250	Oct 10, 2038	U-3950			
	12138250	Oct 10, 2038	U-3951			
<u>SELUMETINIB SULFATE - KOSELUGO</u>						
N 213756 001	11813246	Mar 26, 2029	DP		NCE	Apr 10, 2025
	7425637	Mar 13, 2025	DS		ODE-288	Apr 10, 2027
	8178693	Mar 13, 2025	DS DP			
	9156795	Dec 12, 2026	DS DP			
	9562017	Dec 12, 2026	DS U-2800			
<u>SELUMETINIB SULFATE - KOSELUGO</u>						
N 213756 002	11813246	Mar 26, 2029	DP		NCE	Apr 10, 2025
	7425637	Mar 13, 2025	DS		ODE-288	Apr 10, 2027
	8178693	Mar 13, 2025	DS DP			
	9156795	Dec 12, 2026	DS DP			
	9562017	Dec 12, 2026	DS U-2800			
<u>SEMAGLUTIDE - OZEMPIC</u>						
N 209637 001	10220155	Jul 17, 2026	DP			
	10335462	Jun 21, 2033	U-2580			
	10357616	Jan 20, 2026	DP			
	10376652	Jan 20, 2026	DP			
	11097063	Jul 17, 2026	DP			
	11311679	Jan 20, 2026	DP			
	11446443	Oct 20, 2025	DP			
	8114833	Aug 13, 2025	DP			
	8129343	Dec 05, 2031	DS DP U-2202			
	8536122	Mar 20, 2026	DS DP U-2202			
	8684969	Oct 20, 2025	DP			
	8920383	Jul 17, 2026	DP			
	9108002	Jan 20, 2026	DP			
	9132239	Feb 01, 2032	DP			
	9457154	Sep 27, 2027	DP			
	9616180	Jan 20, 2026	DP			
	9687611	Feb 27, 2027	DP			
	9775953	Jul 17, 2026	DP			
	9861757	Jan 20, 2026	DP			
	RE46363	Aug 03, 2026	DP			
<u>SEMAGLUTIDE - OZEMPIC</u>						
N 209637 002	10220155	Jul 17, 2026	DP			
	10335462	Jun 21, 2033	U-2580			
	10357616	Jan 20, 2026	DP			
	10376652	Jan 20, 2026	DP			
	11097063	Jul 17, 2026	DP			
	11311679	Jan 20, 2026	DP			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>SEMAGLUTIDE - OZEMPIC</u>						
N 209637 002	11446443	Oct 20, 2025	DP			
	8114833	Aug 13, 2025	DP			
	8129343	Dec 05, 2031	DS DP U-2202			
	8536122	Mar 20, 2026	DS DP U-2202			
	8684969	Oct 20, 2025	DP			
	8920383	Jul 17, 2026	DP			
	9108002	Jan 20, 2026	DP			
	9132239	Feb 01, 2032	DP			
	9457154	Sep 29, 2027	DP			
	9616180	Jan 20, 2026	DP			
	9687611	Feb 27, 2027	DP			
	9775953	Jul 17, 2026	DP			
	9861757	Jan 20, 2026	DP			
	RE46363	Aug 03, 2026	DP			
<u>SEMAGLUTIDE - OZEMPIC</u>						
N 209637 003	10220155	Jul 17, 2026	DP		D-185	Mar 28, 2025
	10357616	Jan 20, 2026	DP			
	10376652	Jan 20, 2026	DP			
	11097063	Jul 17, 2026	DP			
	11311679	Jan 20, 2026	DP			
	11446443	Oct 20, 2025	DP			
	8114833	Aug 13, 2025	DP			
	8129343	Dec 05, 2031	DS DP U-3355			
	8536122	Mar 20, 2026	DS DP U-3355			
	8684969	Oct 20, 2025	DP			
	8920383	Jul 17, 2026	DP			
	9108002	Jan 20, 2026	DP			
	9132239	Feb 01, 2032	DP			
	9457154	Sep 29, 2027	DP			
	9616180	Jan 20, 2026	DP			
	9687611	Feb 27, 2027	DP			
	9775953	Jul 17, 2026	DP			
	9861757	Jan 20, 2026	DP			
	RE46363	Aug 03, 2026	DP			
<u>SEMAGLUTIDE - OZEMPIC</u>						
N 209637 004	10220155	Jul 17, 2026	DP			
	10335462	Jun 21, 2033	DP U-2580			
	10357616	Jan 20, 2026	DP			
	10376652	Jan 20, 2026	DP			
	11097063	Jul 17, 2026	DP			
	11311679	Jan 20, 2026	DP			
	11446443	Oct 20, 2025	DP			
	8114833	Aug 13, 2025	DP			
	8129343	Dec 05, 2031	DS DP U-3469			
	8536122	Mar 20, 2026	DS DP U-3469			
	8684969	Oct 20, 2025	DP			
	8920383	Jul 17, 2026	DP			
	9108002	Jan 20, 2026	DP			
	9132239	Feb 01, 2032	DP			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>SEMAGLUTIDE - OZEMPIC</u>						
N 209637	004 9457154	Sep 29, 2027	DP			
	9616180	Jan 20, 2026	DP			
	9687611	Feb 27, 2027	DP			
	9775953	Jul 17, 2026	DP			
	9861757	Jan 20, 2026	DP			
	RE46363	Aug 03, 2026	DP			
<u>SEMAGLUTIDE - RYBELSUS</u>						
N 213051	001 10086047	Dec 16, 2031	DP			
	10278923	May 02, 2034			U-2628	
	10933120	Mar 15, 2033	DP			
	10960052	Dec 16, 2031	DP			
	11382957	Dec 16, 2031	DP			
	11759501	Mar 15, 2033	DP			
	11759502	Mar 15, 2033	DP			
	11759503	Mar 15, 2033	DP			
	8129343	Dec 05, 2031	DS DP		U-2628	
	8536122	Mar 20, 2026	DS DP		U-2628	
	9278123	Dec 16, 2031	DP		U-2628	
<u>SEMAGLUTIDE - RYBELSUS</u>						
N 213051	002 10086047	Dec 16, 2031	DP			
	10278923	May 02, 2034			U-2628	
	10933120	Mar 15, 2033	DP			
	10960052	Dec 16, 2031	DP			
	11382957	Dec 16, 2031	DP			
	11759501	Mar 15, 2033	DP			
	11759502	Mar 15, 2033	DP			
	11759503	Mar 15, 2033	DP			
	8129343	Dec 05, 2031	DS DP		U-2628	
	8536122	Mar 20, 2026	DS DP		U-2628	
	9278123	Dec 16, 2031	DP		U-2628	
<u>SEMAGLUTIDE - RYBELSUS</u>						
N 213051	003 10086047	Dec 16, 2031	DP			
	10278923	May 02, 2034			U-2628	
	10933120	Mar 15, 2033	DP			
	10960052	Dec 16, 2031	DP			
	11382957	Dec 16, 2031	DP			
	11759501	Mar 15, 2033	DP			
	11759502	Mar 15, 2033	DP			
	11759503	Mar 15, 2033	DP			
	8129343	Dec 05, 2031	DS DP		U-2628	
	8536122	Mar 20, 2026	DS DP		U-2628	
	9278123	Dec 16, 2031	DP		U-2628	
<u>SEMAGLUTIDE - RYBELSUS</u>						
N 213051	004 11833248	Feb 01, 2039	DP			U-2628
<u>SEMAGLUTIDE - RYBELSUS</u>						
N 213051	005 11833248	Feb 01, 2039	DP			U-2628

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>SEMAGLUTIDE - RYBELSUS</u>						
N 213051	006 11833248	Feb 01, 2039	DP U-2628			
<u>SEMAGLUTIDE - WEGOVY</u>						
N 215256	001 10888605	Aug 24, 2038	DP U-3162		I-935	Mar 08, 2027
	11318191	Feb 17, 2041	DP U-3162		NPP	Dec 23, 2025
	11752198	Aug 24, 2038	DP U-3162			
	8129343	Dec 05, 2031	DS DP			
	8536122	Mar 20, 2026	DS DP			
	9764003	Jun 21, 2033	U-3161			
<u>SEMAGLUTIDE - WEGOVY</u>						
N 215256	002 10888605	Aug 24, 2038	DP U-3162		I-935	Mar 08, 2027
	11318191	Feb 17, 2041	DP U-3162		NPP	Dec 23, 2025
	11752198	Aug 24, 2038	DP U-3162			
	8129343	Dec 05, 2031	DS DP			
	8536122	Mar 20, 2026	DS DP			
	9764003	Jun 21, 2033	U-3161			
<u>SEMAGLUTIDE - WEGOVY</u>						
N 215256	003 10888605	Aug 24, 2038	DP U-3162		I-935	Mar 08, 2027
	11318191	Feb 17, 2041	DP U-3162		NPP	Dec 23, 2025
	11752198	Aug 24, 2038	DP U-3162			
	8129343	Dec 05, 2031	DS DP			
	8536122	Mar 20, 2026	DS DP			
	9764003	Jun 21, 2033	U-3161			
<u>SEMAGLUTIDE - WEGOVY</u>						
N 215256	004 10888605	Aug 24, 2038	DP U-3162		D-190	Jul 21, 2026
	11318191	Feb 17, 2041	DP U-3162		I-935	Mar 08, 2027
	11752198	Aug 24, 2038	DP U-3162		NPP	Dec 23, 2025
	8129343	Dec 05, 2031	DS DP			
	8536122	Mar 20, 2026	DS DP			
	9764003	Jun 21, 2033	U-3161			
<u>SEMAGLUTIDE - WEGOVY</u>						
N 215256	005 10888605	Aug 24, 2038	DP U-3162		I-935	Mar 08, 2027
	11318191	Feb 17, 2041	DP U-3162		NPP	Dec 23, 2025
	11752198	Aug 24, 2038	DP U-3162			
	12029779	Oct 10, 2038	U-3162			
	8129343	Dec 05, 2031	DS DP			
	8536122	Mar 20, 2026	DS DP			
	9764003	Jun 21, 2033	U-3161			
<u>SETMELANOTIDE ACETATE - IMCIVREE</u>						
N 213793	001 11129869	Jul 04, 2034	DP		I-892	Jun 16, 2025
	8039435	Oct 13, 2027	DS DP		NCE	Nov 25, 2025
	9458195	Oct 13, 2027	DS DP		ODE-336	Nov 25, 2027
					ODE-402	Jun 16, 2029
<u>SEVELAMER CARBONATE - RENVELA</u>						
N 022127	001 7985418	Oct 27, 2025	DP			
<u>SEVELAMER CARBONATE - RENVELA</u>						
N 022318	001 9095509	Dec 06, 2030	DP			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>SEVELAMER CARBONATE - RENVELA</u>						
N 022318	002 9095509	Dec 06, 2030	DP			
<u>SILDENAFIL CITRATE - REVATIO</u>						
N 021845	001				M-287	Jan 31, 2026
					ODE-469	Jan 31, 2030
<u>SILDENAFIL CITRATE - REVATIO</u>						
N 022473	001				M-287	Jan 31, 2026
<u>SILDENAFIL CITRATE - REVATIO</u>						
N 203109	001				M-287	Jan 31, 2026
<u>SILDENAFIL CITRATE - LIOREV</u>						
N 214952	001 11337979	Dec 24, 2038	DP U-3582			
	11464778	Dec 24, 2038	DP U-3582			
	11759468	Dec 24, 2038	DP U-3582			
	12005062	Dec 24, 2038	DP U-3582			
<u>SIMEPREVIR SODIUM - OLYSIO</u>						
N 205123	001 7671032	May 19, 2025	DS DP			
	8148399	Sep 05, 2029	DS DP U-1467			
	8349869	Jul 28, 2026	DS DP U-1467			
	8741926	Jul 28, 2026	DS U-1467			
	8754106	Jul 28, 2026	DS U-1467			
	9040562	Jul 28, 2026	DS DP U-1467			
	9353103	Jul 28, 2026	U-1467			
	9623022	Jul 28, 2026	U-1467			
	9856265	Jul 28, 2026	DS DP U-1467			
<u>SIMVASTATIN - FLOLIPID</u>						
N 206679	001 10300041	Apr 26, 2027	DP			
	9597289	Feb 23, 2030	DP			
<u>SIMVASTATIN - FLOLIPID</u>						
N 206679	002 10300041	Apr 26, 2027	DP			
	9597289	Feb 23, 2030	DP			
<u>SIMVASTATIN; SITAGLIPTIN PHOSPHATE - JUVISYNC</u>						
N 202343	001 7326708	Apr 11, 2026	DS DP U-1188			
	7326708*PED	Oct 11, 2026				
<u>SIMVASTATIN; SITAGLIPTIN PHOSPHATE - JUVISYNC</u>						
N 202343	002 7326708	Apr 11, 2026	DS DP U-1188			
	7326708*PED	Oct 11, 2026				
<u>SIMVASTATIN; SITAGLIPTIN PHOSPHATE - JUVISYNC</u>						
N 202343	003 7326708	Apr 11, 2026	DS DP U-1188			
	7326708*PED	Oct 11, 2026				
<u>SIMVASTATIN; SITAGLIPTIN PHOSPHATE - JUVISYNC</u>						
N 202343	004 7326708	Apr 11, 2026	DS DP U-1188			
	7326708*PED	Oct 11, 2026				
<u>SIMVASTATIN; SITAGLIPTIN PHOSPHATE - JUVISYNC</u>						
N 202343	005 7326708	Apr 11, 2026	DS DP U-1188			
	7326708*PED	Oct 11, 2026				

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>SIMVASTATIN; SITAGLIPTIN PHOSPHATE - JUVISYNC</u>						
N 202343	006 7326708	Apr 11, 2026	DS DP U-1188			
	7326708*PED	Oct 11, 2026				
<u>SINCALIDE - SINCALIDE</u>						
N 210850	001 11110063	Apr 20, 2038	DP U-3481			
	11110063	Apr 20, 2038	DP U-3482			
	11110063	Apr 20, 2038	DP U-3483			
	11318100	Apr 20, 2038	DP U-3477			
	11318100	Apr 20, 2038	DP U-3478			
	11318100	Apr 20, 2038	DP U-3479			
	11318100	Apr 20, 2038	DP U-3480			
	11737983	Apr 20, 2038	DP U-3482			
	11737983	Apr 20, 2038	DP U-3689			
<u>SINECATECHINS - VEREGEN</u>						
N 021902	001 7858662	Oct 02, 2026	DP U-172			
	9770406	Jul 12, 2025	DP U-172			
<u>SIPONIMOD - MAYZENT</u>						
N 209884	001 11944602	Jul 24, 2036	U-3889		M-274	Mar 01, 2025
	12071402	Jan 05, 2032	DP			
	7939519	Aug 27, 2028	DS DP			
	8492441	Nov 30, 2030	U-2511			
<u>SIPONIMOD - MAYZENT</u>						
N 209884	002 11944602	Jul 24, 2036	U-3889		M-274	Mar 01, 2025
	12071402	Jan 05, 2032	DP			
	7939519	Aug 27, 2028	DS DP			
	8492441	Nov 30, 2030	U-2511			
<u>SIPONIMOD - MAYZENT</u>						
N 209884	003 11944602	Jul 24, 2036	U-3889		M-274	Mar 01, 2025
	12071402	Jan 05, 2032	DP			
	7939519	Aug 27, 2028	DS DP			
	8492441	Nov 30, 2030	U-2511			
<u>SIROLIMUS - FYARRO</u>						
N 213312	001 10206887	Apr 15, 2030	DP		ODE-386	Nov 22, 2028
	10705070	Mar 05, 2036	DP			
	10973806	Jun 29, 2036	U-3258			
	11497737	Oct 28, 2040	DP			
	12061183	Mar 05, 2036	DP			
	12133844	Jun 29, 2036	U-4040			
	12133844	Jun 29, 2036	U-4041			
	12133844	Jun 29, 2036	U-4042			
	8911786	Feb 14, 2029	DP U-3259			
<u>SIROLIMUS - HYFTOR</u>						
N 213478	001				NP	Mar 22, 2025
					ODE-391	Mar 22, 2029
<u>SITAGLIPTIN - ZITUVIO</u>						
N 211566	001 10925871	Feb 25, 2035	DP			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>SITAGLIPTIN - ZITUVIO</u>						
N 211566	002 10925871	Feb 25, 2035	DP			
<u>SITAGLIPTIN - ZITUVIO</u>						
N 211566	003 10925871	Feb 25, 2035	DP			
<u>SITAGLIPTIN PHOSPHATE - JANUVIA</u>						
N 021995	001 7326708	Nov 24, 2026	DS DP U-802			
	7326708*PED	May 24, 2027				
<u>SITAGLIPTIN PHOSPHATE - JANUVIA</u>						
N 021995	002 7326708	Nov 24, 2026	DS DP U-802			
	7326708*PED	May 24, 2027				
<u>SITAGLIPTIN PHOSPHATE - JANUVIA</u>						
N 021995	003 7326708	Nov 24, 2026	DS DP U-802			
	7326708*PED	May 24, 2027				
<u>SODIUM NITRITE - SODIUM NITRITE</u>						
N 203922	001 8568793	Dec 24, 2031	DS DP			
	9687506	Feb 10, 2030	DP U-3394			
	9687506	Feb 10, 2030	DP U-3395			
<u>SODIUM NITRITE; SODIUM THIOSULFATE - NITHIODOLE</u>						
N 201444	001 10479686	Jul 07, 2030	DP U-3390			
	11753301	Feb 10, 2030	DS DP U-3681			
	8496973	Mar 29, 2031	DS DP U-1419			
	8568793	Dec 24, 2031	DS DP			
	9345724	Jul 07, 2030	DS DP U-2015			
	9585912	Jul 07, 2030	DS DP			
	9687506	Feb 10, 2030	DP U-3394			
	9687506	Feb 10, 2030	DP U-3395			
<u>SODIUM OXYBATE - XYREM</u>						
N 021196	001 10213400	Mar 15, 2033	U-2499		ODE-231	Oct 26, 2025
	10213400*PED	Sep 15, 2033			PED	Apr 26, 2026
	10864181	Mar 15, 2033	U-1532			
	10864181*PED	Sep 15, 2033				
	11253494	Mar 15, 2033	U-3323			
	11253494	Mar 15, 2033	U-3324			
	11253494*PED	Sep 15, 2033				
	11986446	Mar 15, 2033	U-3324			
	8772306	Mar 15, 2033	U-1532			
	8772306*PED	Sep 15, 2033				
	9050302	Mar 15, 2033	U-1532			
	9050302*PED	Sep 15, 2033				
	9486426	Mar 15, 2033	U-1532			
	9486426*PED	Sep 15, 2033				
<u>SODIUM OXYBATE - LUMRYZ</u>						
N 214755	001 10272062	Jul 21, 2037	DP		NP	May 01, 2026
	10736866	Jul 21, 2037	DP		ODE-431	May 01, 2030
	10925844	Feb 28, 2040	DP		ODE-494	Oct 16, 2031
	10952986	Jul 21, 2037	U-3601			
	10973795	Jul 21, 2037	DP			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>SODIUM OXYBATE - LUMRYZ</u>						
N 214755 001	11000498	Jul 21, 2037	DP U-3580			
	11052061	Jul 21, 2037	DP			
	11065224	Jul 21, 2037	DP			
	11400065	Jul 21, 2037	U-3579			
	11504347	Jul 21, 2037	DP			
	11583510	Feb 07, 2042	U-3578			
	11602512	Jul 21, 2037	U-3577			
	11602513	Jul 21, 2037	U-3576			
	11766418	Jul 21, 2037	DP			
	11779557	Mar 16, 2042	U-3705			
	11839597	Jul 21, 2037	DP			
	11896572	Jul 21, 2037	DP			
	11986451	Jul 21, 2037	U-3934			
	12097175	Jul 21, 2037	DP			
	12097176	Jul 21, 2037	U-4006			
	12109186	Jul 21, 2037	DP			
	12115142	Jul 21, 2037	DP			
	12115143	Jul 21, 2037	DP			
	12115144	Jul 21, 2037	DP			
	12115145	Jul 21, 2037	DP			
	12128021	Jul 21, 2037	U-4022			
	12138239	Jul 21, 2037	DP			
	12144793	Jul 21, 2037	DP			
<u>SODIUM OXYBATE - LUMRYZ</u>						
N 214755 002	10272062	Jul 21, 2037	DP		NP	May 01, 2026
	10736866	Jul 21, 2037	DP		ODE-431	May 01, 2030
	10925844	Feb 28, 2040	DP		ODE-494	Oct 16, 2031
	10952986	Jul 21, 2037	U-3601			
	10973795	Jul 21, 2037	DP			
	11000498	Jul 21, 2037	DP U-3580			
	11052061	Jul 21, 2037	DP			
	11065224	Jul 21, 2037	DP			
	11400065	Jul 21, 2037	U-3579			
	11504347	Jul 21, 2037	DP			
	11583510	Feb 07, 2042	U-3578			
	11602512	Jul 21, 2037	U-3577			
	11602513	Jul 21, 2037	U-3576			
	11766418	Jul 21, 2037	DP			
	11779557	Mar 16, 2042	U-3705			
	11826335	Jul 21, 2037	U-3751			
	11839597	Jul 21, 2037	DP			
	11896572	Jul 21, 2037	DP			
	11986451	Jul 21, 2037	U-3934			
	12097175	Jul 21, 2037	DP			
	12097176	Jul 21, 2037	U-4006			
	12109186	Jul 21, 2037	DP			
	12115142	Jul 21, 2037	DP			
	12115143	Jul 21, 2037	DP			
	12115144	Jul 21, 2037	DP			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>SODIUM OXYBATE - LUMRYZ</u>						
N 214755 002	12115145	Jul 21, 2037	DP			
	12128021	Jul 21, 2037	U-4022			
	12138239	Jul 21, 2037	DP			
	12144793	Jul 21, 2037	DP			
<u>SODIUM OXYBATE - LUMRYZ</u>						
N 214755 003	10272062	Jul 21, 2037	DP		NP	May 01, 2026
	10736866	Jul 21, 2037	DP		ODE-431	May 01, 2030
	10925844	Feb 28, 2040	DP		ODE-494	Oct 16, 2031
	10952986	Jul 21, 2037	U-3601			
	10973795	Jul 21, 2037	DP			
	11000498	Jul 21, 2037	DP U-3580			
	11052061	Jul 21, 2037	DP			
	11065224	Jul 21, 2037	DP			
	11400065	Jul 21, 2037	U-3579			
	11504347	Jul 21, 2037	DP			
	11583510	Feb 07, 2042	U-3578			
	11602512	Jul 21, 2037	U-3577			
	11602513	Jul 21, 2037	U-3576			
	11766418	Jul 21, 2037	DP			
	11779557	Mar 16, 2042	U-3705			
	11839597	Jul 21, 2037	DP			
	11896572	Jul 21, 2037	DP			
	11986451	Jul 21, 2037	U-3934			
	12097175	Jul 21, 2037	DP			
	12097176	Jul 21, 2037	U-4006			
	12109186	Jul 21, 2037	DP			
	12115142	Jul 21, 2037	DP			
	12115143	Jul 21, 2037	DP			
	12115144	Jul 21, 2037	DP			
	12115145	Jul 21, 2037	DP			
	12128021	Jul 21, 2037	U-4022			
	12138239	Jul 21, 2037	DP			
	12144793	Jul 21, 2037	DP			
<u>SODIUM OXYBATE - LUMRYZ</u>						
N 214755 004	10272062	Jul 21, 2037	DP		NP	May 01, 2026
	10736866	Jul 21, 2037	DP		ODE-431	May 01, 2030
	10925844	Feb 28, 2040	DP		ODE-494	Oct 16, 2031
	10952986	Jul 21, 2037	U-3601			
	10973795	Jul 21, 2037	DP			
	11000498	Jul 21, 2037	DP U-3580			
	11052061	Jul 21, 2037	DP			
	11065224	Jul 21, 2037	DP			
	11400065	Jul 21, 2037	U-3579			
	11504347	Jul 21, 2037	DP			
	11583510	Feb 07, 2042	U-3578			
	11602512	Jul 21, 2037	U-3577			
	11602513	Jul 21, 2037	U-3576			
	11766418	Jul 21, 2037	DP			
	11779557	Mar 16, 2042	U-3705			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>SODIUM OXYBATE - LUMRYZ</u>						
N 214755 004	11839597	Jul 21, 2037	DP			
	11896572	Jul 21, 2037	DP			
	11986451	Jul 21, 2037		U-3934		
	12097175	Jul 21, 2037	DP			
	12097176	Jul 21, 2037		U-4006		
	12109186	Jul 21, 2037	DP			
	12115142	Jul 21, 2037	DP			
	12115143	Jul 21, 2037	DP			
	12115144	Jul 21, 2037	DP			
	12115145	Jul 21, 2037	DP			
	12128021	Jul 21, 2037		U-4022		
	12138239	Jul 21, 2037	DP			
	12144793	Jul 21, 2037	DP			
<u>SODIUM PHENYLBUTYRATE - OLPRUVA</u>						
N 214860 001	11154521	Oct 17, 2036	DP			
	11202767	Oct 17, 2036		U-3502		
	11433041	Oct 17, 2036	DP			
<u>SODIUM PHENYLBUTYRATE - OLPRUVA</u>						
N 214860 002	11154521	Oct 17, 2036	DP			
	11202767	Oct 17, 2036		U-3502		
	11433041	Oct 17, 2036	DP			
<u>SODIUM PHENYLBUTYRATE - OLPRUVA</u>						
N 214860 003	11154521	Oct 17, 2036	DP			
	11202767	Oct 17, 2036		U-3502		
	11433041	Oct 17, 2036	DP			
<u>SODIUM PHENYLBUTYRATE - OLPRUVA</u>						
N 214860 004	11154521	Oct 17, 2036	DP			
	11202767	Oct 17, 2036		U-3502		
	11433041	Oct 17, 2036	DP			
<u>SODIUM PHENYLBUTYRATE - OLPRUVA</u>						
N 214860 005	11154521	Oct 17, 2036	DP			
	11202767	Oct 17, 2036		U-3502		
	11433041	Oct 17, 2036	DP			
<u>SODIUM PHENYLBUTYRATE - OLPRUVA</u>						
N 214860 006	11154521	Oct 17, 2036	DP			
	11202767	Oct 17, 2036		U-3502		
	11433041	Oct 17, 2036	DP			
<u>SODIUM PHENYLBUTYRATE; TAURURSODIOL - RELYVRIO</u>						
N 216660 001	10251896	Dec 24, 2033		U-3460	NCE	Sep 29, 2027
	10857162	Dec 24, 2033		U-3460	ODE-411	Sep 29, 2029
	11071742	Dec 24, 2033	DP			
	11583542	Jul 27, 2040	DP			
	9872865	Dec 24, 2033		U-3460		
<u>SODIUM PHOSPHATE, DIBASIC, ANHYDROUS; SODIUM PHOSPHATE, MONOBASIC, MONOHYDRATE - OSMOPREP</u>						
N 021892 001	7687075	Jun 22, 2028	DS DP			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>SODIUM THIOSULFATE - SODIUM THIOSULFATE</u>						
N 203923	001 10479686	Jul 07, 2030	DP U-3390			
	11753301	Feb 10, 2030	DS DP U-3682			
	8496973	Mar 29, 2031	DS DP U-1419			
	9345724	Jul 07, 2030	DS DP U-2015			
	9585912	Jul 07, 2030	DS DP			
<u>SODIUM THIOSULFATE - PEDMARK</u>						
N 212937	001 10596190	Jan 05, 2038	U-3443	Y	NP	Sep 20, 2025
	11291728	Jul 01, 2039	DP		ODE-384	Sep 20, 2029
	11510984	Jul 01, 2039	DP			
	11617793	Jul 01, 2039	DP			
	11964018	Jul 01, 2039	U-3898			
	11992530	Jul 01, 2039	U-3948			
	11998604	Jul 01, 2039	U-3952			
<u>SODIUM ZIRCONIUM CYCLOSILICATE - LOKELMA</u>						
N 207078	001 10300087	Oct 14, 2035	DS U-2312			
	10335432	Feb 10, 2032	U-2312			
	10398730	Feb 10, 2032	U-2312			
	10413569	Feb 10, 2032	DS			
	10695365	Oct 22, 2033	DS			
	11406662	Feb 10, 2032	DS			
	11738044	Oct 14, 2035	U-2312			
	8802152	Apr 19, 2032	DS			
	8808750	Feb 10, 2032	U-2312			
	8877255	Oct 22, 2033	DS			
	9592253	Oct 14, 2035	DS U-2312			
	9844567	Feb 10, 2032	U-2312			
	9861658	Feb 10, 2032	U-2312			
	9913860	Oct 22, 2033	DS U-2312			
<u>SODIUM ZIRCONIUM CYCLOSILICATE - LOKELMA</u>						
N 207078	002 10300087	Oct 14, 2035	DS U-2312			
	10398730	Feb 10, 2032	U-2312			
	10413569	Feb 10, 2032	DS			
	10695365	Oct 22, 2033	DS			
	11406662	Feb 10, 2032	DS			
	11738044	Oct 14, 2035	U-2312			
	8802152	Apr 19, 2032	DS			
	8808750	Feb 10, 2032	U-2312			
	8877255	Oct 22, 2033	DS			
	9592253	Oct 14, 2035	DS U-2312			
	9844567	Feb 10, 2032	U-2312			
	9861658	Feb 10, 2032	U-2312			
	9913860	Oct 22, 2033	DS U-2312			
<u>SOFOSEBUVIR - SOVALDI</u>						
N 204671	001 7964580	Mar 26, 2029	DS DP U-1470		ODE*	Aug 28, 2026
	7964580*PED	Sep 26, 2029				
	8334270	Mar 21, 2028	DS DP U-1470			
	8334270*PED	Sep 21, 2028				
	8580765	Mar 21, 2028	DS DP U-1470			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>SOFOSEBUVIR - SOVALDI</u>						
N 204671	001	8580765*PED	Sep 21, 2028			
		8618076	Dec 11, 2030	DS DP U-1470		
		8618076*PED	Jun 11, 2031			
		8633309	Mar 26, 2029	DS DP U-1470		
		8633309*PED	Sep 26, 2029			
		8889159	Mar 26, 2029	DP U-1470		
		8889159*PED	Sep 26, 2029			
		9085573	Mar 21, 2028	DS DP U-1470		
		9085573*PED	Sep 21, 2028			
		9284342	Sep 13, 2030	DS DP U-1470		
		9284342*PED	Mar 13, 2031			
		9549941	Mar 26, 2029	U-1958		
		9549941*PED	Sep 26, 2029			
<u>SOFOSEBUVIR - SOVALDI</u>						
N 204671	002	7964580	Mar 26, 2029	DS DP U-1470	ODE*	Aug 28, 2026
		7964580*PED	Sep 26, 2029			
		8334270	Mar 21, 2028	DS DP U-1470		
		8334270*PED	Sep 21, 2028			
		8580765	Mar 21, 2028	DS DP U-1470		
		8580765*PED	Sep 21, 2028			
		8618076	Dec 11, 2030	DS DP U-1470		
		8618076*PED	Jun 11, 2031			
		8633309	Mar 26, 2029	DS DP U-1470		
		8633309*PED	Sep 26, 2029			
		8889159	Mar 26, 2029	DP U-1470		
		8889159*PED	Sep 26, 2029			
		9085573	Mar 21, 2028	DS DP U-1470		
		9085573*PED	Sep 21, 2028			
		9284342	Sep 13, 2030	DS DP U-1470		
		9284342*PED	Mar 13, 2031			
<u>SOFOSEBUVIR - SOVALDI</u>						
N 212480	001	7964580	Mar 26, 2029	DS DP U-1470	ODE-258	Aug 28, 2026
		7964580*PED	Sep 26, 2029			
		8334270	Mar 21, 2028	DS DP U-1470		
		8334270*PED	Sep 21, 2028			
		8580765	Mar 21, 2028	DS DP U-1470		
		8580765*PED	Sep 21, 2028			
		8618076	Dec 11, 2030	DS DP U-1470		
		8618076*PED	Jun 11, 2031			
		8633309	Mar 26, 2029	DS DP U-1470		
		8633309*PED	Sep 26, 2029			
		8889159	Mar 26, 2029	DP U-1470		
		8889159*PED	Sep 26, 2029			
		9085573	Mar 21, 2028	DS DP U-1470		
		9085573*PED	Sep 21, 2028			
		9284342	Sep 13, 2030	DS DP U-1470		
		9284342*PED	Mar 13, 2031			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>SOFOSBUVIR - SOVALDI</u>						
N 212480 002	7964580	Mar 26, 2029	DS DP U-1470		ODE-258	Aug 28, 2026
	7964580*PED	Sep 26, 2029				
	8334270	Mar 21, 2028	DS DP U-1470			
	8334270*PED	Sep 21, 2028				
	8580765	Mar 21, 2028	DS DP U-1470			
	8580765*PED	Sep 21, 2028				
	8618076	Dec 11, 2030	DS DP U-1470			
	8618076*PED	Jun 11, 2031				
	8633309	Mar 26, 2029	DS DP U-1470			
	8633309*PED	Sep 26, 2029				
	8889159	Mar 26, 2029	DP U-1470			
	8889159*PED	Sep 26, 2029				
	9085573	Mar 21, 2028	DS DP U-1470			
	9085573*PED	Sep 21, 2028				
	9284342	Sep 13, 2030	DS DP U-1470			
	9284342*PED	Mar 13, 2031				
<u>SOFOSBUVIR; VELPATASVIR - EPCLUSA</u>						
N 208341 001	10086011	Jan 30, 2034	U-1470		M-277	Apr 27, 2025
	10086011*PED	Jul 30, 2034			ODE-293	Mar 19, 2027
	11116783	Jan 30, 2034	DP U-1470		ODE-376	Jun 10, 2028
	11116783*PED	Jul 30, 2034			PED	Sep 19, 2027
	7964580	Mar 26, 2029	DS DP U-1470		PED	Dec 10, 2028
	7964580*PED	Sep 26, 2029				
	8334270	Mar 21, 2028	DS DP U-1470			
	8334270*PED	Sep 21, 2028				
	8575135	Nov 16, 2032	DS DP U-1470			
	8575135*PED	May 16, 2033				
	8580765	Mar 21, 2028	DS DP U-1470			
	8580765*PED	Sep 21, 2028				
	8618076	Dec 11, 2030	DS DP U-1470			
	8618076*PED	Jun 11, 2031				
	8633309	Mar 26, 2029	DS DP U-1470			
	8633309*PED	Sep 26, 2029				
	8735372	Mar 21, 2028	U-1470			
	8735372*PED	Sep 21, 2028				
	8889159	Mar 26, 2029	DP U-1470			
	8889159*PED	Sep 26, 2029				
	8921341	Nov 16, 2032	DS DP U-1470			
	8921341*PED	May 16, 2033				
	8940718	Nov 16, 2032	DS DP U-1470			
	8940718*PED	May 16, 2033				
	9085573	Mar 21, 2028	DS DP U-1470			
	9085573*PED	Sep 21, 2028				
	9284342	Sep 13, 2030	DS DP U-1470			
	9284342*PED	Mar 13, 2031				
	9757406	Jan 30, 2034	DP			
	9757406*PED	Jul 30, 2034				

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>SOFOSBUVIR; VELPATASVIR - EPCLUSA</u>						
N 208341 002	10086011	Jan 30, 2034	U-1470		M-277	Apr 27, 2025
	10086011*PED	Jul 30, 2034			ODE-293	Mar 19, 2027
	11116783	Jan 30, 2034	DP U-1470		ODE-376	Jun 10, 2028
	11116783*PED	Jul 30, 2034			PED	Sep 19, 2027
	7964580	Mar 26, 2029	DS DP U-1470		PED	Dec 10, 2028
	7964580*PED	Sep 26, 2029				
	8334270	Mar 21, 2028	DS DP U-1470			
	8334270*PED	Sep 21, 2028				
	8575135	Nov 16, 2032	DS DP U-1470			
	8575135*PED	May 16, 2033				
	8580765	Mar 21, 2028	DS DP U-1470			
	8580765*PED	Sep 21, 2028				
	8618076	Dec 11, 2030	DS DP U-1470			
	8618076*PED	Jun 11, 2031				
	8633309	Mar 26, 2029	DS DP U-1470			
	8633309*PED	Sep 26, 2029				
	8735372	Mar 21, 2028	U-1470			
	8735372*PED	Sep 21, 2028				
	8889159	Mar 26, 2029	DP U-1470			
	8889159*PED	Sep 26, 2029				
	8921341	Nov 16, 2032	DS DP U-1470			
	8921341*PED	May 16, 2033				
	8940718	Nov 16, 2032	DS DP U-1470			
	8940718*PED	May 16, 2033				
	9085573	Mar 21, 2028	DS DP U-1470			
	9085573*PED	Sep 21, 2028				
	9284342	Sep 13, 2030	DS DP U-1470			
	9284342*PED	Mar 13, 2031				
	9757406	Jan 30, 2034	DP			
	9757406*PED	Jul 30, 2034				
<u>SOFOSBUVIR; VELPATASVIR - EPCLUSA</u>						
N 214187 001	11116783	Jan 30, 2034	DP U-1470		M-277	Apr 27, 2025
	11116783*PED	Jul 30, 2034			ODE-376	Jun 10, 2028
	11707479	Jan 30, 2034	DP U-1470		PED	Dec 10, 2028
	11707479*PED	Jul 30, 2034				
	7964580	Mar 26, 2029	DS DP U-1470			
	7964580*PED	Sep 26, 2029				
	8334270	Mar 21, 2028	DS DP U-1470			
	8334270*PED	Sep 21, 2028				
	8575135	Nov 16, 2032	DS DP U-1470			
	8575135*PED	May 16, 2033				
	8580765	Mar 21, 2028	DS DP U-1470			
	8580765*PED	Sep 21, 2028				
	8618076	Dec 11, 2030	DS DP U-1470			
	8618076*PED	Jun 11, 2031				
	8633309	Mar 26, 2029	DS DP U-1470			
	8633309*PED	Sep 26, 2029				
	8735372	Mar 21, 2028	U-1470			
	8735372*PED	Sep 21, 2028				

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>SOFOBUVIR; VELPATASVIR - EPCLUSA</u>						
N 214187 001	8889159	Mar 26, 2029	DP U-1470			
	8889159*PED	Sep 26, 2029				
	8921341	Nov 16, 2032	DS DP U-1470			
	8921341*PED	May 16, 2033				
	8940718	Nov 16, 2032	DS DP U-1470			
	8940718*PED	May 16, 2033				
	9085573	Mar 21, 2028	DS DP U-1470			
	9085573*PED	Sep 21, 2028				
	9284342	Sep 13, 2030	DS DP U-1470			
	9284342*PED	Mar 13, 2031				
<u>SOFOBUVIR; VELPATASVIR - EPCLUSA</u>						
N 214187 002	11116783	Jan 30, 2034	DP U-1470		M-277	Apr 27, 2025
	11116783*PED	Jul 30, 2034			ODE-376	Jun 10, 2028
	11707479	Jan 30, 2034	DP U-1470		PED	Dec 10, 2028
	11707479*PED	Jul 30, 2034				
	7964580	Mar 26, 2029	DS DP U-1470			
	7964580*PED	Sep 26, 2029				
	8334270	Mar 21, 2028	DS DP U-1470			
	8334270*PED	Sep 21, 2028				
	8575135	Nov 16, 2032	DS DP U-1470			
	8575135*PED	May 16, 2033				
	8580765	Mar 21, 2028	DS DP U-1470			
	8580765*PED	Sep 21, 2028				
	8618076	Dec 11, 2030	DS DP U-1470			
	8618076*PED	Jun 11, 2031				
	8633309	Mar 26, 2029	DS DP U-1470			
	8633309*PED	Sep 26, 2029				
	8735372	Mar 21, 2028	U-1470			
	8735372*PED	Sep 21, 2028				
	8889159	Mar 26, 2029	DP U-1470			
	8889159*PED	Sep 26, 2029				
	8921341	Nov 16, 2032	DS DP U-1470			
	8921341*PED	May 16, 2033				
	8940718	Nov 16, 2032	DS DP U-1470			
	8940718*PED	May 16, 2033				
	9085573	Mar 21, 2028	DS DP U-1470			
	9085573*PED	Sep 21, 2028				
	9284342	Sep 13, 2030	DS DP U-1470			
	9284342*PED	Mar 13, 2031				
<u>SOFOBUVIR; VELPATASVIR; VOXILAPREVIR - VOSEVI</u>						
N 209195 001	10912814	Jun 01, 2037	DP			
	11116783	Jan 30, 2034	DP U-2039			
	11116783	Jan 30, 2034	DP U-2040			
	11116783*PED	Jul 30, 2034				
	11338007	Jun 01, 2037	DP U-2039			
	11338007	Jun 01, 2037	DP U-2040			
	11338007*PED	Dec 01, 2037				
	7964580	Mar 26, 2029	DS DP U-2039			
	7964580	Mar 26, 2029	DS DP U-2040			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>SOFOSBUVIR; VELPATASVIR; VOXILAPREVIR - VOSEVI</u>						
N 209195 001	7964580*PED	Sep 26, 2029				
	8334270	Mar 21, 2028	DS DP U-2039			
	8334270	Mar 21, 2028	DS DP U-2040			
	8334270*PED	Sep 21, 2028				
	8575135	Nov 16, 2032	DS DP U-2039			
	8575135	Nov 16, 2032	DS DP U-2040			
	8575135*PED	May 16, 2033				
	8580765	Mar 21, 2028	DS DP U-2039			
	8580765	Mar 21, 2028	DS DP U-2040			
	8580765*PED	Sep 21, 2028				
	8618076	Dec 11, 2030	DS DP U-2039			
	8618076	Dec 11, 2030	DS DP U-2040			
	8618076*PED	Jun 11, 2031				
	8633309	Mar 26, 2029	DS DP U-2039			
	8633309	Mar 26, 2029	DS DP U-2040			
	8633309*PED	Sep 26, 2029				
	8735372	Mar 21, 2028	DS DP U-2039			
	8735372	Mar 21, 2028	DS DP U-2040			
	8735372*PED	Sep 21, 2028				
	8889159	Mar 26, 2029	DS DP U-2039			
	8889159	Mar 26, 2029	DS DP U-2040			
	8889159*PED	Sep 26, 2029				
	8921341	Nov 16, 2032	DS DP U-2039			
	8921341	Nov 16, 2032	DS DP U-2040			
	8921341*PED	May 16, 2033				
	8940718	Nov 16, 2032	DS DP U-2039			
	8940718	Nov 16, 2032	DS DP U-2040			
	8940718*PED	May 16, 2033				
	8957046	Mar 21, 2028	U-2039			
	8957046	Mar 21, 2028	U-2040			
	9085573	Mar 21, 2028	DS DP U-2039			
	9085573	Mar 21, 2028	DS DP U-2040			
	9085573*PED	Sep 21, 2028				
	9284342	Sep 13, 2030	DS DP U-2039			
	9284342	Sep 13, 2030	DS DP U-2040			
	9284342*PED	Mar 13, 2031				
	9296782	Jul 17, 2034	DS DP			
	9868745	Nov 16, 2032	DS DP			
<u>SOPPIRONIUM BROMIDE - SOFDRA</u>						
N 217347 001	10383846	Mar 14, 2034	U-2398		NCE	Jun 20, 2029
	10947192	May 22, 2034	U-2398			
	10952990	May 22, 2034	U-2398			
	10959983	May 22, 2034	U-2398			
	10961191	May 22, 2034	U-2398			
	11026919	May 22, 2034	DP U-2398			
	11034652	May 22, 2034	DP U-2398			
	11052067	May 22, 2034	DP U-2398			
	11084788	May 22, 2034	DP U-2398			
	11123325	Jul 20, 2037	DP U-2398			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>SOPFIRONIUM BROMIDE - SOFDRA</u>						
N 217347	001 11566000	May 22, 2040	DS DP			
	11584715	May 22, 2040	DS DP			
	8147809	Mar 26, 2027	DS DP			
	8628759	Nov 13, 2026		U-2398		
	9220707	Mar 14, 2034		U-2398		
	9492429	Mar 14, 2034	DP	U-2398		
	9895350	Mar 14, 2034		U-2398		
<u>SOLIFENACIN SUCCINATE - VESICARE LS</u>						
N 209529	001 9918970	May 18, 2031	DP			
<u>SOLRIAMFETOL HYDROCHLORIDE - SUNOSI</u>						
N 211230	001 10195151	Sep 05, 2037	DP		ODE-254	Jun 17, 2026
	10351517	Jun 07, 2026		U-2548		
	10512609	Sep 05, 2037		U-2548		
	10912754	Jun 01, 2038		U-3082		
	10940133	Mar 19, 2040		U-3099		
	10959976	Jun 01, 2038		U-3151		
	11160779	Mar 19, 2040		U-3521		
	11439597	Sep 05, 2037	DP			
	11560354	Mar 06, 2039	DP	U-3520		
	11648232	Jun 01, 2038		U-3602		
	11753368	Jun 07, 2026		U-2548		
	11771666	Dec 30, 2042		U-3693		
	11771667	Dec 30, 2042		U-3693		
	11779554	Dec 30, 2042		U-3693		
	11793776	Dec 30, 2042		U-3693		
	11839598	Mar 19, 2040		U-3765		
	11839599	Mar 19, 2040		U-3764		
	11850226	Mar 19, 2040		U-3775		
	11850227	Mar 19, 2040		U-3775		
	11850228	Mar 19, 2040		U-3775		
	11857528	Mar 19, 2040		U-3521		
	11865098	Jun 01, 2038		U-2548		
	11872203	Dec 30, 2042		U-3693		
	11872204	Dec 30, 2042		U-3693		
	11969404	Mar 19, 2040		U-3892		
	11986454	Mar 19, 2040		U-3775		
	11986455	Mar 19, 2040		U-3521		
	11998639	Sep 05, 2037	DP	U-2548		
	12005036	Dec 30, 2042		U-3693		
	12036194	Dec 30, 2042		U-3693		
	12064411	Dec 30, 2042		U-3693		
	12090126	Dec 30, 2042		U-3693		
	12102609	Dec 30, 2042		U-3693		
	8440715	Jun 11, 2031		U-2548		
	8877806	Jun 07, 2026		U-2548		
	9604917	Jun 07, 2026		U-2548		
<u>SOLRIAMFETOL HYDROCHLORIDE - SUNOSI</u>						
N 211230	002 10195151	Sep 05, 2037	DP		ODE-254	Jun 17, 2026

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>SOLRIAMFETOL HYDROCHLORIDE - SUNOSI</u>						
N 211230 002	10351517	Jun 07, 2026	U-2548			
	10512609	Sep 05, 2037	U-2548			
	10912754	Jun 01, 2038	U-3082			
	10959976	Jun 01, 2038	U-3151			
	11160779	Mar 19, 2040	U-3521			
	11439597	Sep 05, 2037	DP			
	11560354	Mar 06, 2039	DP U-3520			
	11648232	Jun 01, 2038	U-3602			
	11753368	Jun 07, 2026	U-2548			
	11771666	Dec 30, 2042	U-3693			
	11771667	Dec 30, 2042	U-3693			
	11779554	Dec 30, 2042	U-3693			
	11793776	Dec 30, 2042	U-3693			
	11839598	Mar 19, 2040	U-3765			
	11839599	Mar 19, 2040	U-3764			
	11850226	Mar 19, 2040	U-3775			
	11850227	Mar 19, 2040	U-3775			
	11850228	Mar 19, 2040	U-3775			
	11857528	Mar 19, 2040	U-3521			
	11865098	Jun 01, 2038	U-2548			
	11872203	Dec 30, 2042	U-3693			
	11872204	Dec 30, 2042	U-3693			
	11969404	Mar 19, 2040	U-3892			
	11986454	Mar 19, 2040	U-3775			
	11986455	Mar 19, 2040	U-3521			
	11998639	Sep 05, 2037	DP U-2548			
	12005036	Dec 30, 2042	U-3693			
	12036194	Dec 30, 2042	U-3693			
	12064411	Dec 30, 2042	U-3693			
	12090126	Dec 30, 2042	U-3693			
	12102609	Dec 30, 2042	U-3693			
	8440715	Jun 11, 2031	U-2548			
	8877806	Jun 07, 2026	U-2548			
	9604917	Jun 07, 2026	U-2548			
<u>SONIDEGIB PHOSPHATE - ODOMZO</u>						
N 205266 001	10266523	Mar 30, 2036	DS DP			
	8063043	Sep 15, 2029	DS DP			
	8178563	Jul 24, 2029	DS U-1722			
<u>SORAFENIB TOSYLATE - NEXAVAR</u>						
N 021923 001	8877933	Dec 24, 2027	DS DP U-1624			
	9737488	Sep 10, 2028	DP U-1480			
	9737488	Sep 10, 2028	DP U-1696			
	9737488	Sep 10, 2028	DP U-2107			
<u>SOTAGLIFLOZIN - INPEFA</u>						
N 216203 001	7781577	May 04, 2028	DS DP U-3628		NCE	May 26, 2028
	8217156	Oct 07, 2030	DS DP			
	8476413	May 29, 2028	DS DP U-3628			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>SOTAGLIFLOZIN - INPEFA</u>						
N 216203 002	7781577	May 04, 2028	DS DP U-3628		NCE	May 26, 2028
	8217156	Oct 07, 2030	DS DP			
	8476413	May 29, 2028	DS DP U-3628			
<u>SOTALOL HYDROCHLORIDE - SOTALOL HYDROCHLORIDE</u>						
N 022306 001	10512620	Aug 14, 2038	U-2769			
	10512620	Aug 14, 2038	U-3547			
	10799138	Apr 05, 2039	U-3125			
	10799138	Apr 05, 2039	U-3549			
	11583216	Aug 21, 2039	U-3549			
	11696902	Aug 14, 2038	U-2769			
<u>SOTALOL HYDROCHLORIDE - SOTYLIZE</u>						
N 205108 001	10206895	Apr 01, 2034	DP U-2096			
	10206895	Apr 01, 2034	DP U-2494			
	11013703	Apr 01, 2034	DP			
	11850222	Nov 19, 2034	U-2096			
	11850222	Nov 19, 2034	U-2494			
	9724297	Aug 31, 2035	DP U-2096			
<u>SOTORASIB - LUMAKRAS</u>						
N 214665 001	10519146	May 21, 2038	DS DP		NCE	May 28, 2026
	11236091	May 20, 2040	DS DP U-3306		ODE-352	May 28, 2028
	11426404	Sep 15, 2040	U-3306			
	11827635	May 20, 2040	DS DP U-3306			
<u>SOTORASIB - LUMAKRAS</u>						
N 214665 002	10519146	May 21, 2038	DS DP		NCE	May 28, 2026
	11236091	May 20, 2040	DS DP U-3306		ODE-352	May 28, 2028
	11426404	Sep 15, 2040	U-3306			
	11827635	May 20, 2040	DS DP U-3306			
<u>SOTORASIB - LUMAKRAS</u>						
N 214665 003	10519146	May 21, 2038	DS DP		NCE	May 28, 2026
	11236091	May 20, 2040	DS DP U-3306		ODE*	May 28, 2028
	11426404	Sep 15, 2040	U-3306			
	11827635	May 20, 2040	DS DP U-3306			
<u>SPARSENTAN - FILSPARI</u>						
N 216403 001	9993461	Mar 29, 2030	U-3269		NCE	Feb 17, 2028
	9993461	Mar 29, 2030	U-3993		ODE-389	Feb 17, 2030
					ODE-493	Sep 05, 2031
<u>SPARSENTAN - FILSPARI</u>						
N 216403 002	9993461	Mar 29, 2030	U-3269		NCE	Feb 17, 2028
	9993461	Mar 29, 2030	U-3993		ODE-389	Feb 17, 2030
					ODE-493	Sep 05, 2031
<u>SPINOSAD - NATROBA</u>						
N 022408 001	9895388	Nov 25, 2033	U-3365			
<u>SPIRONOLACTONE - CAROSPIR</u>						
N 209478 001	10493083	Oct 28, 2036	DP			
	10624906	Oct 28, 2036	DP			
	10660907	Oct 28, 2036	DP			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>SPIRONOLACTONE - CAROSPIR</u>						
N 209478	001 10888570	Oct 28, 2036	DP			
	11389461	Oct 28, 2036	DP			
	11395828	Oct 28, 2036		U-2109		
	11395828	Oct 28, 2036		U-3401		
	11395828	Oct 28, 2036		U-3402		
	11491166	Oct 28, 2036	DP			
	9757394	Oct 28, 2036	DP	U-2109		
<u>STIRIPENTOL - DIACOMIT</u>						
N 206709	001				M-281	Jul 14, 2025
					ODE-198	Aug 20, 2025
					ODE-403	Jul 14, 2029
<u>STIRIPENTOL - DIACOMIT</u>						
N 206709	002				M-281	Jul 14, 2025
					ODE-198	Aug 20, 2025
					ODE-403	Jul 14, 2029
<u>STIRIPENTOL - DIACOMIT</u>						
N 207223	001				NPP	Jul 14, 2025
					ODE-198	Aug 20, 2025
					ODE-403	Jul 14, 2029
<u>STIRIPENTOL - DIACOMIT</u>						
N 207223	002				NPP	Jul 14, 2025
					ODE-198	Aug 20, 2025
					ODE-403	Jul 14, 2029
<u>SUFENTANIL CITRATE - DSUVIA</u>						
N 209128	001 10245228	Jan 05, 2027	DP	U-1351		
	10342762	Jan 05, 2027	DP			
	10507180	Jan 05, 2027	DP	U-1351		
	10896751	Mar 16, 2030	DP			
	11672738	Feb 02, 2038	DP			
	11676691	Mar 16, 2030	DP			
	12033733	Mar 16, 2030	DP			
	8202535	Oct 22, 2030		U-1351		
	8226978	Jan 05, 2027	DP	U-1351		
	8231900	Jan 05, 2027	DP			
	8252328	Jan 05, 2027	DP			
	8252329	Jan 05, 2027	DP			
	8535714	Jan 05, 2027	DP	U-1351		
	8574189	Mar 16, 2030	DP			
	8778393	Jan 05, 2027		U-1351		
	8778394	Jan 05, 2027		U-1351		
	8865211	Jan 05, 2027		U-1351		
	8865743	Oct 22, 2030		U-1351		
	8945592	Jul 29, 2031	DP			
	9320710	Jan 05, 2027		U-1351		
	9744129	Jan 05, 2027	DP			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>SUGAMMADEX SODIUM - BRIDION</u>						
N 022225	001	RE44733	Jan 27, 2026	DS DP	U-1794	
		RE44733*PED	Jul 27, 2026			
<u>SUGAMMADEX SODIUM - BRIDION</u>						
N 022225	002	RE44733	Jan 27, 2026	DS DP	U-1794	
		RE44733*PED	Jul 27, 2026			
<u>SULFUR HEXAFLUORIDE LIPID-TYPE A MICROSOPHERES - LUMASON</u>						
N 203684	001	10232061	Jul 06, 2038	DP		
		10335502	Jul 06, 2038	DP		
		11723869	May 15, 2039	DP	U-3666	
		11723869	May 15, 2039	DP	U-3667	
		11723869	May 15, 2039	DP	U-3668	
<u>SUMATRIPTAN - TOSYMRA</u>						
N 210884	001	10603305	Jun 16, 2030	DP	U-1719	
		11337962	Jun 16, 2030	DP	U-1719	
		12090139	Jun 16, 2030	DP		
		8268791	May 09, 2026	DP		
		8440631	May 09, 2026	DP	U-1719	
		9211282	Jul 19, 2031	DP	U-1719	
		9283280	May 09, 2026	DP		
		9610280	Jun 16, 2030	DP	U-1719	
		9974770	Jun 16, 2030	DP	U-1719	
<u>SUMATRIPTAN SUCCINATE - SUMAVEL DOSEPRO</u>						
N 022239	001	7776007	Nov 22, 2026	DP		
		7901385	Jul 31, 2026	DP		
<u>SUMATRIPTAN SUCCINATE - ALSUMA</u>						
N 022377	001	7811254	Aug 26, 2027	DP	U-1083	
<u>SUMATRIPTAN SUCCINATE - ZECUITY</u>						
N 202278	001	7973058	Apr 12, 2027		U-1328	
		8155737	Apr 12, 2027		U-1328	
		8366600	Apr 21, 2029		U-1327	
		8470853	Apr 12, 2027		U-1328	
		8597272	Apr 12, 2027	DP		
		8983594	Nov 19, 2030	DP	U-1328	
		9272137	Sep 07, 2027	DP		
		9327114	Oct 08, 2032	DP	U-1328	
		9427578	Apr 12, 2027	DP	U-1328	
<u>SUMATRIPTAN SUCCINATE - ONZETRA XSAIL</u>						
N 206099	001	10076614	Oct 20, 2034	DP		
		10076615	Jul 30, 2029		U-2010	
		10076615	Jul 30, 2029		U-2011	
		10076615	Jul 30, 2029		U-2404	
		10124132	Mar 06, 2027	DP	U-1719	
		10124132	Mar 06, 2027	DP	U-2010	
		10124132	Mar 06, 2027	DP	U-2011	
		10398859	Dec 19, 2027	DP		
		10478574	Nov 04, 2033		U-2404	
		10722667	Dec 30, 2028	DP		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>SUMATRIPTAN SUCCINATE - ONZETRA XSAIL</u>						
N 206099	001	11571531	Feb 23, 2026			
				U-1809		
		7975690	Aug 18, 2025	DP	U-1809	
		8550073	Oct 22, 2029	DP		
		8590530	Sep 15, 2025	DP	U-1809	
		8875704	Apr 07, 2028	DP	U-1809	
		8899229	Aug 18, 2030	DP		
		8978647	Dec 06, 2030	DP		
		9108015	Sep 15, 2025	DP		
		9649456	Oct 21, 2030	DP	U-1719	
		9649456	Oct 21, 2030	DP	U-2010	
		9649456	Oct 21, 2030	DP	U-2011	
<u>SUMATRIPTAN SUCCINATE - ZEMBRACE SYMTOUCH</u>						
N 208223	001	10537554	Jan 29, 2036			
				U-72		
		11364224	Jan 29, 2036			
				U-72		
		12097183	Jan 29, 2036	DP		
<u>SUVOREXANT - BELSOMRA</u>						
N 204569	001	10098892	May 29, 2033	DP		
		11980623	May 29, 2033	DP		
		7951797	Nov 20, 2029	DS DP	U-620	
<u>SUVOREXANT - BELSOMRA</u>						
N 204569	002	10098892	May 29, 2033	DP		
		11980623	May 29, 2033	DP		
		7951797	Nov 20, 2029	DS DP	U-620	
<u>SUVOREXANT - BELSOMRA</u>						
N 204569	003	10098892	May 29, 2033	DP		
		11980623	May 29, 2033	DP		
		7951797	Nov 20, 2029	DS DP	U-620	
<u>SUVOREXANT - BELSOMRA</u>						
N 204569	004	10098892	May 29, 2033	DP		
		11980623	May 29, 2033	DP		
		7951797	Nov 20, 2029	DS DP	U-620	
<u>TACROLIMUS - PROGRAF</u>						
N 050708	001				ODE-294	May 24, 2025
					ODE-360	Jul 16, 2028
<u>TACROLIMUS - PROGRAF</u>						
N 050708	002				ODE-294	May 24, 2025
					ODE-360	Jul 16, 2028
<u>TACROLIMUS - PROGRAF</u>						
N 050708	003				ODE-294	May 24, 2025
					ODE-360	Jul 16, 2028
<u>TACROLIMUS - PROGRAF</u>						
N 050709	001				ODE-294	May 24, 2025
					ODE-360	Jul 16, 2028
<u>TACROLIMUS - ASTAGRAF XL</u>						
N 204096	001				ODE*	May 24, 2025

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>TACROLIMUS - ASTAGRAF XL</u>						
N 204096	002				ODE*	May 24, 2025
<u>TACROLIMUS - ASTAGRAF XL</u>						
N 204096	003				ODE*	May 24, 2025
<u>TACROLIMUS - ENVARSUS XR</u>						
N 206406	001	10166190	May 30, 2028	DP		
		10864199	May 30, 2028	U-2677		
		10864199	May 30, 2028	U-2678		
		11110081	May 30, 2028	U-2678		
		11123331	May 30, 2028	U-2677		
		11419823	May 30, 2028	DP		
		12083103	May 30, 2028	U-2678		
		8664239	Aug 30, 2028	U-1752		
		8664239	Aug 30, 2028	U-2677		
		8664239	Aug 30, 2028	U-2678		
		8685998	Aug 30, 2028	DP U-1752		
		8685998	Aug 30, 2028	DP U-2677		
		8685998	Aug 30, 2028	DP U-2678		
		9549918	May 30, 2028	DP		
<u>TACROLIMUS - ENVARSUS XR</u>						
N 206406	002	10166190	May 30, 2028	DP		
		10864199	May 30, 2028	U-2677		
		10864199	May 30, 2028	U-2678		
		11110081	May 30, 2028	U-2678		
		11123331	May 30, 2028	U-2677		
		11419823	May 30, 2028	DP		
		12083103	May 30, 2028	U-2678		
		8664239	Aug 30, 2028	U-1752		
		8664239	Aug 30, 2028	U-2677		
		8664239	Aug 30, 2028	U-2678		
		8685998	Aug 30, 2028	DP U-1752		
		8685998	Aug 30, 2028	DP U-2677		
		8685998	Aug 30, 2028	DP U-2678		
		9549918	May 30, 2028	DP		
<u>TACROLIMUS - ENVARSUS XR</u>						
N 206406	003	10166190	May 30, 2028	DP		
		10864199	May 30, 2028	U-2677		
		10864199	May 30, 2028	U-2678		
		11110081	May 30, 2028	U-2678		
		11123331	May 30, 2028	U-2677		
		11419823	May 30, 2028	DP		
		12083103	May 30, 2028	U-2678		
		8664239	Aug 30, 2028	U-1752		
		8664239	Aug 30, 2028	U-2677		
		8664239	Aug 30, 2028	U-2678		
		8685998	Aug 30, 2028	DP U-1752		
		8685998	Aug 30, 2028	DP U-2677		
		8685998	Aug 30, 2028	DP U-2678		
		9549918	May 30, 2028	DP		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>TACROLIMUS - ENVARSUS XR</u>						
N 206406 003	10166190	May 30, 2028	DP			
	10864199	May 30, 2028		U-2677		
	10864199	May 30, 2028		U-2678		
	11110081	May 30, 2028		U-2678		
	11123331	May 30, 2028		U-2677		
	11419823	May 30, 2028	DP			
	12083103	May 30, 2028		U-2678		
	8664239	Aug 30, 2028		U-1752		
	8664239	Aug 30, 2028		U-2677		
	8664239	Aug 30, 2028		U-2678		
	8685998	Aug 30, 2028	DP	U-1752		
	8685998	Aug 30, 2028	DP	U-2677		
	8685998	Aug 30, 2028	DP	U-2678		
	9549918	May 30, 2028	DP			
<u>TACROLIMUS - PROGRAF</u>						
N 210115 001					ODE-269	May 24, 2025
					ODE-360	Jul 16, 2028
<u>TACROLIMUS - PROGRAF</u>						
N 210115 002					ODE-269	May 24, 2025
					ODE-360	Jul 16, 2028
<u>TADALAFIL - TADLIO</u>						
N 214522 001	11382917	Dec 24, 2038	DP	U-3397		
	11666576	Dec 24, 2038	DP	U-3397		
	11975006	Dec 24, 2038	DP	U-3397		
<u>TAFAMIDIS - VYNDAMAX</u>						
N 212161 001	7214695	Dec 19, 2025	DS DP		ODE-237	May 03, 2026
	7214696	Dec 19, 2025		U-2524		
	9770441	Aug 31, 2035	DS DP	U-2524		
<u>TAFAMIDIS MEGLUMINE - VYNDAOEL</u>						
N 211996 001	7214695	Dec 19, 2025	DS DP		ODE-237	May 03, 2026
	7214696	Dec 19, 2025		U-2524		
<u>TAFENOQUINE SUCCINATE - ARAKODA</u>						
N 210607 001	10342791	Dec 02, 2035		U-2582		
	10888558	Dec 02, 2035		U-2582		
	11744828	Dec 02, 2035		U-2582		
<u>TAFENOQUINE SUCCINATE - KRINTAFEL</u>						
N 210795 001					ODE-201	Jul 20, 2025
<u>TAFLUPROST - ZIOPTAN</u>						
N 202514 001	10864159	May 28, 2029	DP	U-778		
	9999593	May 28, 2029	DP			
<u>TALAZOPARIB TOSYLATE - TALZENNA</u>						
N 211651 001	10189837	Oct 20, 2031	DS DP		I-920	Jun 20, 2026
	10780088	Jul 27, 2029		U-3651		
	8012976	Oct 19, 2029	DS DP			
	8420650	Jul 27, 2029	DS DP			
	8735392	Oct 20, 2031	DS DP			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>TALAZOPARIB TOSYLATE - TALZENNA</u>						
N 211651	001 9820985	Jul 27, 2029	U-2437			
<u>TALAZOPARIB TOSYLATE - TALZENNA</u>						
N 211651	002 10189837	Oct 20, 2031	DS DP			
	8012976	Oct 19, 2029	DS DP			
	8420650	Jul 27, 2029	DS DP			
	8735392	Oct 20, 2031	DS DP			
	9820985	Jul 27, 2029	U-2437			
<u>TALAZOPARIB TOSYLATE - TALZENNA</u>						
N 211651	003 10189837	Oct 20, 2031	DS DP		I-920	Jun 20, 2026
	10780088	Jul 27, 2029	U-3651			
	8012976	Oct 19, 2029	DS DP			
	8420650	Jul 27, 2029	DS DP			
	8735392	Oct 20, 2031	DS DP			
	9820985	Jul 27, 2029	U-2437			
<u>TALAZOPARIB TOSYLATE - TALZENNA</u>						
N 211651	004 10189837	Oct 20, 2031	DS DP			
	8012976	Oct 19, 2029	DS DP			
	8420650	Jul 27, 2029	DS DP			
	8735392	Oct 20, 2031	DS DP			
	9820985	Jul 27, 2029	U-2437			
<u>TALAZOPARIB TOSYLATE - TALZENNA</u>						
N 211651	005 10189837	Oct 20, 2031	DS DP		I-920	Jun 20, 2026
	10780088	Jul 27, 2029	U-3651			
	8012976	Oct 19, 2029	DS DP			
	8420650	Jul 27, 2029	DS DP			
	8735392	Oct 20, 2031	DS DP			
	9820985	Jul 27, 2029	U-2437			
<u>TALAZOPARIB TOSYLATE - TALZENNA</u>						
N 211651	006 10189837	Oct 20, 2031	DS DP			
	10780088	Jul 27, 2029	U-3651			
	8012976	Oct 19, 2029	DS DP			
	8420650	Jul 27, 2029	DS DP			
	8735392	Oct 20, 2031	DS DP			
	9820985	Jul 27, 2029	U-2437			
<u>TALAZOPARIB TOSYLATE - TALZENNA</u>						
N 217439	001 10189837	Oct 20, 2031	DS DP		I-920	Jun 20, 2026
	10780088	Jul 27, 2029	U-3651			
	8012976	Oct 19, 2029	DS DP			
	8420650	Jul 27, 2029	DS DP			
	8735392	Oct 20, 2031	DS DP			
	9820985	Jul 27, 2029	U-2437			
<u>TALAZOPARIB TOSYLATE - TALZENNA</u>						
N 217439	002 10189837	Oct 20, 2031	DS DP		I-920	Jun 20, 2026
	10780088	Jul 27, 2029	U-3651			
	8012976	Oct 19, 2029	DS DP			
	8420650	Jul 27, 2029	DS DP			
	8735392	Oct 20, 2031	DS DP			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>TALAZOPARIB TOSYLATE - TALZENNA</u>						
N 217439	002 9820985	Jul 27, 2029	U-2437			
<u>TALAZOPARIB TOSYLATE - TALZENNA</u>						
N 217439	003 10189837	Oct 20, 2031	DS DP			
	10780088	Jul 27, 2029	U-3651			
	8012976	Oct 19, 2029	DS DP			
	8420650	Jul 27, 2029	DS DP			
	8735392	Oct 20, 2031	DS DP			
	9820985	Jul 27, 2029	U-2437			
<u>TALAZOPARIB TOSYLATE - TALZENNA</u>						
N 217439	004 10189837	Oct 20, 2031	DS DP		I-920	Jun 20, 2026
	10780088	Jul 27, 2029	U-3651			
	8012976	Oct 19, 2029	DS DP			
	8420650	Jul 27, 2029	DS DP			
	8735392	Oct 20, 2031	DS DP			
	9820985	Jul 27, 2029	U-2437			
<u>TALAZOPARIB TOSYLATE - TALZENNA</u>						
N 217439	005 10189837	Oct 20, 2031	DS DP			
	10780088	Jul 27, 2029	U-3651			
	8012976	Oct 19, 2029	DS DP			
	8420650	Jul 27, 2029	DS DP			
	8735392	Oct 20, 2031	DS DP			
	9820985	Jul 27, 2029	U-2437			
<u>TALAZOPARIB TOSYLATE - TALZENNA</u>						
N 217439	006 10189837	Oct 20, 2031	DS DP			
	10780088	Jul 27, 2029	U-3651			
	8012976	Oct 19, 2029	DS DP			
	8420650	Jul 27, 2029	DS DP			
	8735392	Oct 20, 2031	DS DP			
	9820985	Jul 27, 2029	U-2437			
<u>TAPENTADOL HYDROCHLORIDE - NUCYNTA</u>						
N 022304	001 7994364	Jun 27, 2025	DS DP U-931		NPP	Jul 03, 2026
					PED	Jan 03, 2027
<u>TAPENTADOL HYDROCHLORIDE - NUCYNTA</u>						
N 022304	002 7994364	Jun 27, 2025	DS DP U-931		NPP	Jul 03, 2026
					PED	Jan 03, 2027
<u>TAPENTADOL HYDROCHLORIDE - NUCYNTA</u>						
N 022304	003 7994364	Jun 27, 2025	DS DP U-931		NPP	Jul 03, 2026
					PED	Jan 03, 2027
<u>TAPENTADOL HYDROCHLORIDE - NUCYNTA ER</u>						
N 200533	001 11344512	Apr 21, 2028	U-3391			
	11344512	Apr 21, 2028	U-3392			
	11344512*PED	Oct 21, 2028				
	7994364	Jun 27, 2025	DS DP U-1178			
	7994364	Jun 27, 2025	DS DP U-1276			
	8536130	Sep 22, 2028	U-1276			
	8536130*PED	Mar 22, 2029				

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>TAPENTADOL HYDROCHLORIDE - NUCYNTA ER</u>						
N 200533	002	11344512	Apr 21, 2028	U-3391		
		11344512	Apr 21, 2028	U-3392		
		11344512*PED	Oct 21, 2028			
		7994364	Jun 27, 2025	DS DP U-1178		
		7994364	Jun 27, 2025	DS DP U-1276		
		8536130	Sep 22, 2028	U-1276		
		8536130*PED	Mar 22, 2029			
<u>TAPENTADOL HYDROCHLORIDE - NUCYNTA ER</u>						
N 200533	003	11344512	Apr 21, 2028	U-3391		
		11344512	Apr 21, 2028	U-3392		
		11344512*PED	Oct 21, 2028			
		7994364	Jun 27, 2025	DS DP U-1178		
		7994364	Jun 27, 2025	DS DP U-1276		
		8536130	Sep 22, 2028	U-1276		
		8536130*PED	Mar 22, 2029			
<u>TAPENTADOL HYDROCHLORIDE - NUCYNTA ER</u>						
N 200533	004	11344512	Apr 21, 2028	U-3391		
		11344512	Apr 21, 2028	U-3392		
		11344512*PED	Oct 21, 2028			
		7994364	Jun 27, 2025	DS DP U-1178		
		7994364	Jun 27, 2025	DS DP U-1276		
		8536130	Sep 22, 2028	U-1276		
		8536130*PED	Mar 22, 2029			
<u>TAPENTADOL HYDROCHLORIDE - NUCYNTA ER</u>						
N 200533	005	11344512	Apr 21, 2028	U-3391		
		11344512	Apr 21, 2028	U-3392		
		11344512*PED	Oct 21, 2028			
		7994364	Jun 27, 2025	DS DP U-1178		
		7994364	Jun 27, 2025	DS DP U-1276		
		8536130	Sep 22, 2028	U-1276		
		8536130*PED	Mar 22, 2029			
<u>TAPENTADOL HYDROCHLORIDE - NUCYNTA</u>						
N 203794	001	7994364	Jun 27, 2025	DS DP U-1289	NPP	Jul 03, 2026
					PED	Jan 03, 2027
<u>TAPINAROF - VTAMA</u>						
N 215272	001	10195160	May 19, 2036	DP	NCE	May 23, 2027
		10426743	May 19, 2036	U-2625		
		10647649	Nov 13, 2038	DS		
		11458108	May 19, 2036	DP		
		11590088	Nov 13, 2039	U-2625		
		11597692	Nov 13, 2038	DS DP		
		11612573	May 19, 2036	U-2625		
		11617724	May 19, 2036	DP		
		11622945	May 19, 2036	DP		
<u>TASIMELTEON - HETLIOZ</u>						
N 205677	001	10071977	Feb 12, 2035	DS DP	ODE-330	Dec 01, 2027
		10149829	Jan 25, 2033	U-2477		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>TASIMELTEON - HETLIOZ</u>						
N 205677 001	10149829	Jan 25, 2033	U-3006			
	10179119	Aug 29, 2035	U-3003			
	10376487	Jul 27, 2035	U-2615			
	10376487	Jul 27, 2035	U-3007			
	10449176	Jan 25, 2033	U-2149			
	10610510	Jan 25, 2033	U-2805			
	10610510	Jan 25, 2033	U-3009			
	10610511	Oct 10, 2034	U-2615			
	10610511	Oct 10, 2034	U-3007			
	10829465	Feb 12, 2035	DS DP			
	10945988	Jan 25, 2033	U-2149			
	10980770	Jan 25, 2033	U-3106			
	10980770	Jan 25, 2033	U-3107			
	11141400	Oct 10, 2034	U-2615			
	11141400	Oct 10, 2034	U-3007			
	11266622	Aug 29, 2035	U-3003			
	11285129	Jan 25, 2033	U-3342			
	11285129	Jan 25, 2033	U-3343			
	11566011	Feb 12, 2035	DS DP			
	11633377	Jan 25, 2033	U-2149			
	11633377	Jan 25, 2033	U-3003			
	11759446	Feb 21, 2041	U-3003			
	11760740	Feb 12, 2035	DS DP			
	11786502	Oct 10, 2034	U-3007			
	11786502	Oct 10, 2034	U-3739			
	11826339	Jan 25, 2033	U-3342			
	11826339	Jan 25, 2033	U-3343			
	11833130	Jan 25, 2033	U-2149			
	11833130	Jan 25, 2033	U-3003			
	11850229	Jan 25, 2033	U-3342			
	11850229	Jan 25, 2033	U-3343			
	11918556	Apr 07, 2033	U-3342			
	11918556	Apr 07, 2033	U-3343			
	11918557	Jan 25, 2033	U-3003			
	11918557	Jan 25, 2033	U-3865			
	12049457	Feb 12, 2035	DS DP			
	9060995	Jan 25, 2033	U-1710			
	9539234	Jan 25, 2033	U-1934			
	9539234	Jan 25, 2033	U-3004			
	9549913	Jan 25, 2033	U-1486			
	9730910	May 17, 2034	U-2085			
	9730910	May 17, 2034	U-3005			
	9855241	Jan 25, 2033	U-2149			
	RE46604	Jan 25, 2033	U-2147			
<u>TASIMELTEON - HETLIOZ LO</u>						
N 214517 001	10071977	Feb 12, 2035	DS DP		ODE-329	Dec 01, 2027
	10149829	Jan 25, 2033	U-3006			
	10179119	Aug 29, 2035	U-3003			
	10376487	Jul 27, 2035	U-3007			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>TASIMELTEON - HETLIOZ LQ</u>						
N 214517 001	10610510	Jan 25, 2033	U-3009			
	10610511	Oct 10, 2034	U-3007			
	10829465	Feb 12, 2035	DS DP			
	10980770	Jan 25, 2033	U-3106			
	11141400	Oct 10, 2034	U-3007			
	11202770	Dec 11, 2040	DP			
	11266622	Aug 29, 2035	U-3003			
	11285129	Jan 25, 2033	U-3342			
	11566011	Feb 12, 2035	DS DP			
	11633377	Jan 25, 2033	U-3003			
	11759446	Feb 21, 2041	U-3003			
	11760740	Feb 12, 2035	DS DP			
	11786502	Oct 10, 2034	U-3007			
	11826339	Jan 25, 2033	U-3342			
	11833130	Jan 25, 2033	U-3003			
	11850229	Jan 25, 2033	U-3342			
	11918556	Apr 07, 2033	U-3342			
	11918557	Jan 25, 2033	U-3003			
	12049457	Feb 12, 2035	DS DP			
	9539234	Jan 25, 2033	U-3004			
	9730910	May 17, 2034	U-3005			
<u>TAZAROTENE - TAZAROTENE</u>						
A 217075 001					CGT	Mar 08, 2025
<u>TAZAROTENE - FABIOR</u>						
N 202428 001	10568859	Feb 24, 2030	DP U-2760			
	10688071	Feb 24, 2030	DP U-2760			
	8808716	Feb 24, 2030	DP			
<u>TAZAROTENE - ARAZLO</u>						
N 211882 001	11311482	May 11, 2038	U-2368			
	11679116	Jun 06, 2036	DP			
	12128137	May 11, 2038	DP U-2368			
<u>TAZEMETOSTAT HYDROBROMIDE - TAZVERIK</u>						
N 211723 001	10155002	Sep 12, 2031	U-2736		NCE	Jan 23, 2025
	10245269	Apr 11, 2033	U-2737		ODE-299	Jan 23, 2027
	10245269	Apr 11, 2033	U-2851		ODE-314	Jun 18, 2027
	10245269	Apr 11, 2033	U-2854			
	10369155	Oct 16, 2035	U-2736			
	10369155	Oct 16, 2035	U-2852			
	10369155	Oct 16, 2035	U-2853			
	10420775	Sep 12, 2031	U-2736			
	10420775	Sep 12, 2031	U-2852			
	10420775	Sep 12, 2031	U-2853			
	10786511	Dec 19, 2035	DP			
	10821113	Apr 11, 2033	DS DP			
	11052093	Apr 13, 2032	DS DP U-2736			
	11052093	Apr 13, 2032	DS DP U-2852			
	11052093	Apr 13, 2032	DS DP U-2853			
	11491163	Apr 11, 2033	U-2736			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>TAZEMETOSTAT HYDROBROMIDE - TAZVERIK</u>						
N 211723	001	11491163	Apr 11, 2033	U-2852		
		11491163	Apr 11, 2033	U-2853		
		8410088	Jan 23, 2034	DS DP		
		8691507	Sep 12, 2031	U-2852		
		8765732	Sep 12, 2031	U-2852		
		8765732	Sep 12, 2031	U-2853		
		8895245	Sep 12, 2031	U-2852		
		9090562	Sep 12, 2031	DS DP		
		9175331	Sep 12, 2031	U-2852		
		9333217	Sep 12, 2031	U-2852		
		9334527	Sep 12, 2031	U-2852		
		9394283	Apr 11, 2033	DS DP U-2852		
		9394283	Apr 11, 2033	DS DP U-2853		
		9549931	Sep 12, 2031	U-2736		
		9549931	Sep 12, 2031	U-2852		
		9549931	Sep 12, 2031	U-2853		
		9688665	Aug 22, 2034	U-2736		
		9855275	Sep 12, 2031	U-2736		
		9889138	Oct 16, 2035	U-2736		
		9889138	Oct 16, 2035	U-2852		
		9889138	Oct 16, 2035	U-2853		
		9949999	Sep 12, 2031	U-2851		
<u>TECHNETIUM TC-99M LABELED CARBON - TECHNEGAS KIT</u>						
N 022335	001				NP	Sep 29, 2026
<u>TECHNETIUM TC-99M TETROFOSMIN KIT - MYOVIEW 30ML</u>						
N 020372	002	9549999	Mar 10, 2030	DP		
<u>TECHNETIUM TC-99M TILMANOCEPT - LYMPHOSEEK KIT</u>						
N 202207	001	6409990	May 12, 2025	DS		
		9439985	Jan 30, 2029	DP		
<u>TECOVIRIMAT - TPOXX</u>						
N 208627	001	11890270	Aug 08, 2032	U-2346	ODE-200	Jul 13, 2025
		7737168	Sep 04, 2031	U-2346		
		8039504	Jul 23, 2027	DP		
		9339466	Mar 23, 2031	DS DP		
<u>TECOVIRIMAT - TPOXX</u>						
N 214518	001	10576165	Aug 02, 2031	DP		
		7737168	Sep 04, 2031	U-3377		
		8039504	Jul 23, 2027	DP		
		9233097	Aug 02, 2031	DP		
		9907859	Aug 02, 2031	U-3377		
<u>TEDIZOLID PHOSPHATE - SIVEXTRO</u>						
N 205435	001	10065947	Feb 03, 2030	DP		
		10442829	Feb 03, 2030	DS		
		7816379	Jun 20, 2028	DS DP U-2507		
		8420676	Feb 23, 2028	DS DP U-282		
		8426389	Dec 31, 2030	DS DP U-282		
		9624250	Feb 03, 2030	DS DP U-2507		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>TEDIZOLID PHOSPHATE - SIVEXTRO</u>						
N 205435	001 9988406	Feb 03, 2030	DP			
<u>TEDIZOLID PHOSPHATE - SIVEXTRO</u>						
N 205436	001 10065947	Feb 03, 2030	DP			
	10442829	Feb 03, 2030	DS			
	7816379	Jun 20, 2028	DS DP	U-2507		
	8420676	Feb 23, 2028	DS DP	U-282		
	8426389	Dec 31, 2030	DS DP	U-282		
	9624250	Feb 03, 2030	DS DP	U-2507		
	9988406	Feb 03, 2030	DP			
<u>TEDUGLUTIDE - GATTEX KIT</u>						
N 203441	001 7847061	Nov 01, 2025	U-1320	Y	ODE-240	May 16, 2026
	7847061*PED	May 01, 2026				
	9060992	Nov 01, 2025	U-1320	Y		
	9060992*PED	May 01, 2026				
<u>TELAPREVIR - INCIVEK</u>						
N 201917	001 7820671	Feb 25, 2025	DS DP			
	8431615	May 30, 2028	U-1398			
<u>TELAVANCIN HYDROCHLORIDE - VIBATIV</u>						
N 022110	001 7531623	Jan 01, 2027	DS			
<u>TELAVANCIN HYDROCHLORIDE - VIBATIV</u>						
N 022110	002 7531623	Jan 01, 2027	DS			
<u>TELOTTRISTAT ETIPRATE - XERMELO</u>						
N 208794	001 7553840	Dec 11, 2027	DS			
	7709493	Feb 28, 2031	DS	U-1979		
	7968559	Dec 11, 2027		U-1979		
	8193204	Feb 27, 2031	DS			
	8653094	Dec 19, 2028		U-1979		
<u>TEMSIROLIMUS - TORISEL</u>						
N 022088	001 8026276	Jan 20, 2026	DP			
	8791097	May 10, 2032		U-1550		
	8791097	May 10, 2032		U-1551		
	8791097*PED	Nov 10, 2032				
<u>TENAPANOR HYDROCHLORIDE - IBSRELA</u>						
N 211801	001 12016856	Dec 30, 2029	DS DP			
	8541448	Aug 01, 2033	DS DP			
	8969377	Dec 30, 2029	DS DP			
	9006281	May 02, 2030		U-2626		
	9408840	Dec 30, 2029		U-2626		
<u>TENAPANOR HYDROCHLORIDE - XPHOZAH</u>						
N 213931	001 10272079	Apr 10, 2034		U-3736	NP	Oct 17, 2026
	10272079	Apr 10, 2034		U-381		
	10940146	Apr 10, 2034		U-3736		
	10940146	Apr 10, 2034		U-381		
	12016856	Dec 30, 2029	DS DP			
	8541448	Aug 01, 2033	DS DP			
	8969377	Dec 30, 2029	DS DP			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>TENAPANOR HYDROCHLORIDE - XPHOZAH</u>						
N 213931	002	10272079	Apr 10, 2034	U-3736	NP	Oct 17, 2026
		10272079	Apr 10, 2034	U-381		
		10940146	Apr 10, 2034	U-3736		
		10940146	Apr 10, 2034	U-381		
		12016856	Dec 30, 2029	DS DP		
		8541448	Aug 01, 2033	DS DP		
		8969377	Dec 30, 2029	DS DP		
<u>TENAPANOR HYDROCHLORIDE - XPHOZAH</u>						
N 213931	003	10272079	Apr 10, 2034	U-3736	NP	Oct 17, 2026
		10272079	Apr 10, 2034	U-381		
		10940146	Apr 10, 2034	U-3736		
		10940146	Apr 10, 2034	U-381		
		12016856	Dec 30, 2029	DS DP		
		8541448	Aug 01, 2033	DS DP		
		8969377	Dec 30, 2029	DS DP		
<u>TENOFOVIR ALAFENAMIDE FUMARATE - VEMLIDY</u>						
N 208464	001	7390791	Apr 17, 2025	DS DP	NPP	Oct 17, 2025
		7390791*PED	Oct 17, 2025			
		8754065	Aug 15, 2032	DS DP U-1275		
		8754065	Aug 15, 2032	DS DP U-3880		
		8754065	Aug 15, 2032	DS DP U-999		
		8754065*PED	Feb 15, 2033			
		9296769	Aug 15, 2032	DS DP U-1275		
		9296769	Aug 15, 2032	DS DP U-3880		
		9296769	Aug 15, 2032	DS DP U-999		
		9296769*PED	Feb 15, 2033			
<u>TEPOTINIB HYDROCHLORIDE - TEPMETKO</u>						
N 214096	001	8329692	Oct 30, 2029	DS DP	NCE	Feb 03, 2026
		8580781	Mar 19, 2030	DS DP	ODE-325	Feb 03, 2028
		8658643	Jul 04, 2028	U-3077		
		8921357	May 30, 2028	DS DP		
		8927540	Jul 21, 2028	U-3078		
		9062029	Jul 04, 2028	DP		
		9284300	Apr 29, 2028	DP		
		9403799	Jul 04, 2028	U-3077		
<u>TERAZOSIN HYDROCHLORIDE - TEZRULY</u>						
N 218139	001	11224572	Jan 18, 2042	DP U-3990		
		11224572	Jan 18, 2042	DP U-3991		
<u>TERIFLUNOMIDE - AUBAGIO</u>						
N 202992	001	6794410	Sep 12, 2026	U-1285		
		6794410*PED	Mar 12, 2027			
		8802735	Sep 14, 2030	DP		
		8802735*PED	Mar 14, 2031			
		9186346	Feb 04, 2034	U-1786		
		9186346*PED	Aug 04, 2034			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>TERIFLUNOMIDE - AUBAGIO</u>						
N 202992	002 6794410	Sep 12, 2026	U-1285			
	6794410*PED	Mar 12, 2027				
	8802735	Sep 14, 2030	DP			
	8802735*PED	Mar 14, 2031				
	9186346	Feb 04, 2034	U-1786			
	9186346*PED	Aug 04, 2034				
<u>TERIPARATIDE - FORTEO</u>						
N 021318	001 7517334	Mar 25, 2025	DP			
<u>TERLIPRESSIN ACETATE - TERLIVAZ</u>						
N 022231	001 10335452	Apr 05, 2037	U-3711		NCE ODE-406	Sep 14, 2027 Sep 14, 2029
<u>TESTOSTERONE - ANDROGEL</u>						
N 022309	001 8466136	Oct 12, 2026	DP			
	8466137	Oct 12, 2026	U-1103			
	8466138	Oct 12, 2026	U-1103			
	8486925	Oct 12, 2026	DP			
	8729057	Oct 12, 2026	DP			
	8741881	Oct 12, 2026	U-1103			
	8754070	Oct 12, 2026	DP			
	8759329	Oct 12, 2026	DP			
<u>TESTOSTERONE - ANDROGEL</u>						
N 022309	002 8466136	Oct 12, 2026	DP			
	8466137	Oct 12, 2026	U-1103			
	8466138	Oct 12, 2026	U-1103			
	8486925	Oct 12, 2026	DP			
	8729057	Oct 12, 2026	DP			
	8741881	Oct 12, 2026	U-1103			
	8754070	Oct 12, 2026	DP			
	8759329	Oct 12, 2026	DP			
<u>TESTOSTERONE - ANDROGEL</u>						
N 022309	003 8466136	Oct 12, 2026	DP			
	8466137	Oct 12, 2026	U-1103			
	8466138	Oct 12, 2026	U-1103			
	8486925	Oct 12, 2026	DP			
	8729057	Oct 12, 2026	DP			
	8741881	Oct 12, 2026	U-1103			
	8754070	Oct 12, 2026	DP			
	8759329	Oct 12, 2026	DP			
<u>TESTOSTERONE - AXIRON</u>						
N 022504	001 8419307	Feb 26, 2027	U-1386			
	8435944	Sep 27, 2027	U-1390			
	8807861	Feb 26, 2027	DP U-1563			
	8993520	Jun 02, 2026	U-1390			
	9180194	Jun 02, 2026	U-1390			
	9289586	Feb 26, 2027	U-1390			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>TESTOSTERONE - VOGELXO</u>						
N 204399	002	8785426	Feb 11, 2034	DP U-1531		
		9295675	Feb 11, 2034	DP U-1531		
		9662340	Feb 11, 2034	DP U-1531		
<u>TESTOSTERONE - VOGELXO</u>						
N 204399	003	8785426	Feb 11, 2034	DP U-1531		
		9295675	Feb 11, 2034	DP U-1531		
		9662340	Feb 11, 2034	DP U-1531		
<u>TESTOSTERONE - NATESTO</u>						
N 205488	001	11090312	Mar 17, 2034	U-1616		
<u>TESTOSTERONE CYPIONATE - AZMIRO</u>						
N 216318	001	11311554	Mar 25, 2039	DP		
		11642355	Mar 25, 2039	DP		
		12138271	Mar 25, 2039	DP		
<u>TESTOSTERONE ENANTHATE - XYOSTED (AUTOINJECTOR)</u>						
N 209863	001	10238662	Feb 19, 2035	DP U-2418		
		10279131	Jul 31, 2031	DP		
		10357609	Aug 21, 2031	DP		
		10478560	Jan 24, 2026	DP		
		10646495	Aug 30, 2038	DP		
		10821072	Jun 04, 2033	DP U-2418		
		10881798	Feb 11, 2034	DP		
		10905827	Aug 21, 2031	DP		
		10912782	Feb 19, 2035	DP U-2418		
		11160751	Oct 07, 2034	DP U-2418		
		11191908	Oct 18, 2035	DP		
		11446440	Aug 21, 2031	DP		
		11446441	Jan 24, 2026	DP		
		11497753	Mar 19, 2030	DP		
		11813435	Feb 25, 2035	DP		
		11844804	Jun 04, 2033	U-2418		
		8021335	Oct 04, 2026	DP		
		8562564	Jan 24, 2026	DP		
		9180259	Jan 24, 2026	DP		
		9533102	Jan 24, 2026	DP		
		9629959	Jan 24, 2026	DP		
		9744302	Nov 19, 2035	DP		
		9950125	Sep 04, 2036	DP U-2418		
<u>TESTOSTERONE ENANTHATE - XYOSTED (AUTOINJECTOR)</u>						
N 209863	002	10238662	Feb 19, 2035	DP U-2418		
		10279131	Jul 31, 2031	DP		
		10357609	Aug 21, 2031	DP		
		10478560	Jan 24, 2026	DP		
		10646495	Aug 30, 2038	DP		
		10821072	Jun 04, 2033	DP U-2418		
		10881798	Feb 11, 2034	DP		
		10905827	Aug 21, 2031	DP		
		10912782	Feb 19, 2035	DP U-2418		
		11160751	Oct 07, 2034	DP U-2418		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>TESTOSTERONE ENANTHATE - XYOSTED (AUTOINJECTOR)</u>						
N 209863	002	11191908	Oct 18, 2035	DP		
		11446440	Aug 21, 2031	DP		
		11446441	Jan 24, 2026	DP		
		11497753	Mar 19, 2030	DP		
		11813435	Feb 25, 2035	DP		
		11844804	Jun 04, 2033		U-2418	
		8021335	Oct 04, 2026	DP		
		8562564	Jan 24, 2026	DP		
		9180259	Jan 24, 2026	DP		
		9533102	Jan 24, 2026	DP		
		9629959	Jan 24, 2026	DP		
		9744302	Nov 19, 2035	DP		
		9950125	Sep 04, 2036	DP	U-2418	
<u>TESTOSTERONE ENANTHATE - XYOSTED (AUTOINJECTOR)</u>						
N 209863	003	10238662	Feb 19, 2035	DP	U-2418	
		10279131	Jul 31, 2031	DP		
		10357609	Aug 21, 2031	DP		
		10478560	Jan 24, 2026	DP		
		10646495	Aug 30, 2038	DP		
		10821072	Jun 04, 2033	DP	U-2418	
		10881798	Feb 11, 2034	DP		
		10905827	Aug 21, 2031	DP		
		10912782	Feb 19, 2035	DP	U-2418	
		11160751	Oct 07, 2034	DP	U-2418	
		11191908	Oct 18, 2035	DP		
		11446440	Aug 21, 2031	DP		
		11446441	Jan 24, 2026	DP		
		11497753	Mar 19, 2030	DP		
		11813435	Feb 25, 2035	DP		
		11844804	Jun 04, 2033		U-2418	
		8021335	Oct 04, 2026	DP		
		8562564	Jan 24, 2026	DP		
		9180259	Jan 24, 2026	DP		
		9533102	Jan 24, 2026	DP		
		9629959	Jan 24, 2026	DP		
		9744302	Nov 19, 2035	DP		
		9950125	Sep 04, 2036	DP	U-2418	
<u>TESTOSTERONE UNDECANOATE - AVEED</u>						
N 022219	001	7718640	Mar 14, 2027	DP		
		8338395	May 08, 2027		U-1500	
<u>TESTOSTERONE UNDECANOATE - JATENZO</u>						
N 206089	001	10543219	Apr 12, 2030		U-2506	
		10617696	Apr 12, 2030	DS DP		
		11179402	Apr 14, 2026	DS DP		
		11179403	Apr 12, 2030		U-2506	
		11331325	Jan 06, 2027		U-2506	
		11426416	Apr 12, 2030		U-3420	
		11564933	Apr 12, 2039		U-1103	

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>TESTOSTERONE UNDECANOATE - JATENZO</u>						
N 206089 001	8241664	Mar 29, 2029	DP U-2506			
	8492369	Dec 20, 2030	DP U-2506			
	8778916	Apr 12, 2030	DP			
<u>TESTOSTERONE UNDECANOATE - JATENZO</u>						
N 206089 002	10543219	Apr 12, 2030	U-2506			
	10617696	Apr 12, 2030	DS DP			
	11179402	Apr 14, 2026	DS DP			
	11179403	Apr 12, 2030	U-2506			
	11331325	Jan 06, 2027	U-2506			
	11426416	Apr 12, 2030	U-3420			
	11564933	Apr 12, 2039	U-1103			
	8241664	Mar 29, 2029	DP U-2506			
	8492369	Dec 20, 2030	DP U-2506			
	8778916	Apr 12, 2030	DP			
<u>TESTOSTERONE UNDECANOATE - JATENZO</u>						
N 206089 003	10543219	Apr 12, 2030	U-2506			
	10617696	Apr 12, 2030	DS DP			
	11179402	Apr 14, 2026	DS DP			
	11179403	Apr 12, 2030	U-2506			
	11331325	Jan 06, 2027	U-2506			
	11426416	Apr 12, 2030	U-3420			
	11564933	Apr 12, 2039	U-1103			
	8241664	Mar 29, 2029	DP U-2506			
	8492369	Dec 20, 2030	DP U-2506			
	8778916	Apr 12, 2030	DP			
<u>TESTOSTERONE UNDECANOATE - TLANDO</u>						
N 208088 001	10226473	Nov 30, 2030	U-1500		NP	Mar 28, 2025
	10716794	Nov 30, 2030	U-1500			
	11304960	Jan 08, 2029	DP U-1500			
	11311555	Nov 30, 2030	U-1500			
	11364249	Nov 30, 2030	U-1500			
	11433083	Nov 30, 2030	U-1500			
	11464735	Apr 28, 2041	DP U-1500			
	11559530	Nov 28, 2037	U-1500			
	12011503	Oct 16, 2040	DP U-3964			
	12011503	Oct 16, 2040	DP U-3965			
	8778922	Jan 08, 2029	DP U-1500			
	8865695	Jan 08, 2029	U-1500			
	9943527	Nov 30, 2030	U-1500			
	9949985	Nov 30, 2030	U-1500			
<u>TESTOSTERONE UNDECANOATE - KYZATREX</u>						
N 213953 001	10576089	Dec 31, 2030	DP U-2506		NP	Jul 27, 2025
	10576090	Dec 31, 2030	DP			
	11590146	Dec 31, 2030	DP			
	11617758	Mar 15, 2033	DP U-2506			
<u>TESTOSTERONE UNDECANOATE - KYZATREX</u>						
N 213953 002	10576089	Dec 31, 2030	DP U-2506		NP	Jul 27, 2025
	10576090	Dec 31, 2030	DP			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>TESTOSTERONE UNDECANOATE - KYZATREX</u>						
N 213953	002	11590146	Dec 31, 2030	DP		
		11617758	Mar 15, 2033	DP U-2506		
<u>TESTOSTERONE UNDECANOATE - KYZATREX</u>						
N 213953	003	10576089	Dec 31, 2030	DP U-2506	NP	Jul 27, 2025
		10576090	Dec 31, 2030	DP		
		11590146	Dec 31, 2030	DP		
		11617758	Mar 15, 2033	DP U-2506		
<u>THEOPHYLLINE - THEOPHYLLINE</u>						
A 215312	001				CGT	Mar 26, 2025
<u>THIOTEPA - TEPYLUTE</u>						
N 216984	001	11975013	Aug 16, 2041	DP		
<u>TICAGRELOR - BRILINTA</u>						
N 022433	001	10300065	Jan 27, 2036	U-2541	M-283	May 09, 2025
		10300065	Jan 27, 2036	U-2542	PED	Nov 09, 2025
		10300065*PED	Jul 27, 2036			
		8425934	Apr 17, 2030	DP		
		8425934*PED	Oct 17, 2030			
		RE46276*PED	Apr 30, 2025			
<u>TICAGRELOR - BRILINTA</u>						
N 022433	002	10300065	Jan 27, 2036	U-2541	M-283	May 09, 2025
		10300065	Jan 27, 2036	U-2542	PED	Nov 09, 2025
		10300065*PED	Jul 27, 2036			
		8425934	Apr 17, 2030	DP		
		8425934*PED	Oct 17, 2030			
		RE46276*PED	Apr 30, 2025			
<u>TIGECYCLINE - TYGACIL</u>						
N 021821	001	10588975	Mar 13, 2026	DP		
		7879828	Feb 05, 2029	DP		
		8372995	Oct 08, 2030	DP		
		8975242	Oct 24, 2028	DP		
		9254328	Mar 13, 2026	DP		
		9694078	Mar 13, 2026	DP		
<u>TIGECYCLINE - TIGECYCLINE</u>						
N 211158	001	9855335	Apr 07, 2033	DP		
<u>TIOPRONIN - THIOLA</u>						
N 019569	001				ODE-267	Jun 28, 2026
<u>TIOPRONIN - THIOLA EC</u>						
N 211843	001	11458104	Nov 14, 2038	U-3441	ODE*	Jun 28, 2026
<u>TIOPRONIN - THIOLA EC</u>						
N 211843	002	11458104	Nov 14, 2038	U-3441	ODE*	Jun 28, 2026
<u>TIOTROPIUM BROMIDE - SPIRIVA</u>						
N 021395	001	7694676	Mar 12, 2027	DP		
		7694676*PED	Sep 12, 2027			
		8022082	Jan 19, 2026	DP U-1186		
		8022082*PED	Jul 19, 2026			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>TIOTROPIUM BROMIDE - SPIRIVA</u>						
N 021395	001 9010323	Apr 19, 2030	DP			
<u>TIOTROPIUM BROMIDE - SPIRIVA RESPIMAT</u>						
N 021936	001 7284474*PED	Feb 26, 2025				
	7396341	Oct 10, 2026	DP			
	7396341*PED	Apr 10, 2027				
	7837235	Mar 13, 2028	DP			
	7837235*PED	Sep 13, 2028				
	7896264	May 26, 2025	DP			
	7896264*PED	Nov 26, 2025				
	8733341	Oct 16, 2030	DP			
	8733341*PED	Apr 16, 2031				
	9027967	Mar 31, 2027	DP			
	9027967*PED	Sep 30, 2027				
<u>TIOTROPIUM BROMIDE - SPIRIVA RESPIMAT</u>						
N 021936	002 7284474*PED	Feb 26, 2025				
	7396341	Oct 10, 2026	DP			
	7396341*PED	Apr 10, 2027				
	7837235	Mar 13, 2028	DP			
	7837235*PED	Sep 13, 2028				
	7896264	May 26, 2025	DP			
	7896264*PED	Nov 26, 2025				
	8733341	Oct 16, 2030	DP			
	8733341*PED	Apr 16, 2031				
	9027967	Mar 31, 2027	DP			
	9027967*PED	Sep 30, 2027				
<u>TIPIRACIL HYDROCHLORIDE; TRIFLURIDINE - LONSURE</u>						
N 207981	001 10456399	Feb 03, 2037	U-3657		ODE-229	Feb 22, 2026
	10456399	Feb 03, 2037	U-3658			
	10457666	Jun 17, 2034	DS DP			
	10960004	Feb 03, 2037	U-3657			
	10960004	Feb 03, 2037	U-3658			
	9527833	Jun 17, 2034	DS DP			
	9943537	Sep 05, 2034	U-3659			
	RE46284	Sep 22, 2029	U-1751			
	RE46284	Sep 22, 2029	U-2503			
	RE46284	Sep 22, 2029	U-3656			
<u>TIPIRACIL HYDROCHLORIDE; TRIFLURIDINE - LONSURE</u>						
N 207981	002 10456399	Feb 03, 2037	U-3657		ODE-229	Feb 22, 2026
	10456399	Feb 03, 2037	U-3658			
	10457666	Jun 17, 2034	DS DP			
	10960004	Feb 03, 2037	U-3657			
	10960004	Feb 03, 2037	U-3658			
	9527833	Jun 17, 2034	DS DP			
	9943537	Sep 05, 2034	U-3659			
	RE46284	Sep 22, 2029	U-1751			
	RE46284	Sep 22, 2029	U-2503			
	RE46284	Sep 22, 2029	U-3656			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>TIRBANIBULIN - KLISYRI</u>						
N 213189 001	10323001	Dec 28, 2027	DP		D-192	Jun 07, 2027
	10617693	Mar 12, 2038	U-3015		NCE	Dec 14, 2025
	10669236	Sep 07, 2038	DS DP			
	11497750	Mar 12, 2038	U-3015			
	7300931	Feb 06, 2026	DS DP			
	7851470	Feb 02, 2029	DS DP U-3015			
	8236799	Dec 28, 2025	DS DP			
	8980890	Dec 28, 2025	DS DP			
<u>TIRZEPATIDE - MOUNJARO (AUTOINJECTOR)</u>						
N 215866 001	11357820	Jun 14, 2039	DP		NCE	May 13, 2027
	9474780	Jan 05, 2036	DS DP U-3378			
<u>TIRZEPATIDE - MOUNJARO (AUTOINJECTOR)</u>						
N 215866 002	11357820	Jun 14, 2039	DP		NCE	May 13, 2027
	9474780	Jan 05, 2036	DS DP U-3378			
<u>TIRZEPATIDE - MOUNJARO (AUTOINJECTOR)</u>						
N 215866 003	11357820	Jun 14, 2039	DP		NCE	May 13, 2027
	9474780	Jan 05, 2036	DS DP U-3378			
<u>TIRZEPATIDE - MOUNJARO (AUTOINJECTOR)</u>						
N 215866 004	11357820	Jun 14, 2039	DP		NCE	May 13, 2027
	9474780	Jan 05, 2036	DS DP U-3378			
<u>TIRZEPATIDE - MOUNJARO (AUTOINJECTOR)</u>						
N 215866 005	11357820	Jun 14, 2039	DP		NCE	May 13, 2027
	9474780	Jan 05, 2036	DS DP U-3378			
<u>TIRZEPATIDE - MOUNJARO (AUTOINJECTOR)</u>						
N 215866 006	11357820	Jun 14, 2039	DP		NCE	May 13, 2027
	9474780	Jan 05, 2036	DS DP U-3378			
<u>TIRZEPATIDE - MOUNJARO</u>						
N 215866 007	11357820	Jun 14, 2039	DS DP		NCE	May 13, 2027
	9474780	Jan 05, 2036	DS DP U-3378			
<u>TIRZEPATIDE - MOUNJARO</u>						
N 215866 008	11357820	Jun 14, 2039	DS DP		NCE	May 13, 2027
	9474780	Jan 05, 2036	DS DP U-3378			
<u>TIRZEPATIDE - MOUNJARO</u>						
N 215866 009	11357820	Jun 14, 2039	DS DP		NCE	May 13, 2027
	9474780	Jan 05, 2036	DS DP U-3378			
<u>TIRZEPATIDE - MOUNJARO</u>						
N 215866 010	11357820	Jun 14, 2039	DS DP		NCE	May 13, 2027
	9474780	Jan 05, 2036	DS DP U-3378			
<u>TIRZEPATIDE - MOUNJARO</u>						
N 215866 011	11357820	Jun 14, 2039	DS DP		NCE	May 13, 2027
	9474780	Jan 05, 2036	DS DP U-3378			
<u>TIRZEPATIDE - MOUNJARO</u>						
N 215866 012	11357820	Jun 14, 2039	DS DP		NCE	May 13, 2027
	9474780	Jan 05, 2036	DS DP U-3378			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>TIRZEPATIDE - ZEPBOUND (AUTOINJECTOR)</u>						
N 217806 001	11357820	Jun 14, 2039	DP		M-82	Oct 18, 2027
	11918623	Jun 14, 2039		U-3855	NCE	May 13, 2027
	9474780	Jan 05, 2036	DS DP		NP	Nov 08, 2026
<u>TIRZEPATIDE - ZEPBOUND (AUTOINJECTOR)</u>						
N 217806 002	11357820	Jun 14, 2039	DP		M-82	Oct 18, 2027
	11918623	Jun 14, 2039		U-3855	NCE	May 13, 2027
	9474780	Jan 05, 2036	DS DP		NP	Nov 08, 2026
<u>TIRZEPATIDE - ZEPBOUND (AUTOINJECTOR)</u>						
N 217806 003	11357820	Jun 14, 2039	DP		M-82	Oct 18, 2027
	11918623	Jun 14, 2039		U-3855	NCE	May 13, 2027
	9474780	Jan 05, 2036	DS DP		NP	Nov 08, 2026
<u>TIRZEPATIDE - ZEPBOUND (AUTOINJECTOR)</u>						
N 217806 004	11357820	Jun 14, 2039	DP		M-82	Oct 18, 2027
	11918623	Jun 14, 2039		U-3855	NCE	May 13, 2027
	9474780	Jan 05, 2036	DS DP		NP	Nov 08, 2026
<u>TIRZEPATIDE - ZEPBOUND (AUTOINJECTOR)</u>						
N 217806 005	11357820	Jun 14, 2039	DP		M-82	Oct 18, 2027
	11918623	Jun 14, 2039		U-3855	NCE	May 13, 2027
	9474780	Jan 05, 2036	DS DP		NP	Nov 08, 2026
<u>TIRZEPATIDE - ZEPBOUND (AUTOINJECTOR)</u>						
N 217806 006	11357820	Jun 14, 2039	DP		M-82	Oct 18, 2027
	11918623	Jun 14, 2039		U-3855	NCE	May 13, 2027
	9474780	Jan 05, 2036	DS DP		NP	Nov 08, 2026
<u>TIRZEPATIDE - ZEPBOUND</u>						
N 217806 007	11357820	Jun 14, 2039	DP		M-82	Oct 18, 2027
	11918623	Jun 14, 2039		U-3855	NCE	May 13, 2027
	9474780	Jan 05, 2036	DS DP			
<u>TIRZEPATIDE - ZEPBOUND</u>						
N 217806 008	11357820	Jun 14, 2039	DP		M-82	Oct 18, 2027
	11918623	Jun 14, 2039		U-3855	NCE	May 13, 2027
	9474780	Jan 05, 2036	DS DP			
<u>TIRZEPATIDE - ZEPBOUND</u>						
N 217806 009	11357820	Jun 14, 2039	DP		M-82	Oct 18, 2027
	11918623	Jun 14, 2039		U-3855	NCE	May 13, 2027
	9474780	Jan 05, 2036	DS DP			
<u>TIRZEPATIDE - ZEPBOUND</u>						
N 217806 010	11357820	Jun 14, 2039	DP		M-82	Oct 18, 2027
	11918623	Jun 14, 2039		U-3855	NCE	May 13, 2027
	9474780	Jan 05, 2036	DS DP			
<u>TIRZEPATIDE - ZEPBOUND</u>						
N 217806 011	11357820	Jun 14, 2039	DP		M-82	Oct 18, 2027
	11918623	Jun 14, 2039		U-3855	NCE	May 13, 2027
	9474780	Jan 05, 2036	DS DP			
<u>TIRZEPATIDE - ZEPBOUND</u>						
N 217806 012	11357820	Jun 14, 2039	DP		M-82	Oct 18, 2027

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>TIRZEPATIDE - ZEPBOUND</u>						
N 217806 012	11918623	Jun 14, 2039	U-3855		NCE	May 13, 2027
	9474780	Jan 05, 2036	DS DP			
<u>TIVOZANIB HYDROCHLORIDE - FOTIVDA</u>						
N 212904 001	11504365	Nov 05, 2039	U-3476		NCE	Mar 10, 2026
	6821987	Apr 26, 2025	DS DP U-3100			
<u>TIVOZANIB HYDROCHLORIDE - FOTIVDA</u>						
N 212904 002	11504365	Nov 05, 2039	U-3476		NCE	Mar 10, 2026
	6821987	Apr 26, 2025	DS DP U-3100			
<u>TIZANIDINE HYDROCHLORIDE - ONTRALFY</u>						
N 216190 001	12042484	May 07, 2042	DP U-2779			
<u>TOBRAMYCIN - TOBI PODHALER</u>						
N 201688 001	10207066	Nov 04, 2030	DP U-909			
	7559325	Oct 27, 2025	DP			
	8664187	Jun 20, 2025	U-909			
	8869794	Sep 12, 2028	DP U-909			
<u>TOFACITINIB CITRATE - XELJANZ</u>						
N 203214 001	RE41783	Dec 08, 2025	DS			
<u>TOFACITINIB CITRATE - XELJANZ</u>						
N 203214 002	RE41783	Dec 08, 2025	DS			
<u>TOFACITINIB CITRATE - XELJANZ XR</u>						
N 208246 001	11253523	Mar 14, 2034	U-3326			
	11253523	Mar 14, 2034	U-3327			
	11253523	Mar 14, 2034	U-3328			
	11253523	Mar 14, 2034	U-3329			
	9937181	Mar 14, 2034	DP			
	RE41783	Dec 08, 2025	DS			
<u>TOFACITINIB CITRATE - XELJANZ XR</u>						
N 208246 002	10639309	Mar 14, 2034	DP			
	11253523	Mar 14, 2034	U-3326			
	11253523	Mar 14, 2034	U-3327			
	11253523	Mar 14, 2034	U-3328			
	11253523	Mar 14, 2034	U-3329			
	RE41783	Dec 08, 2025	DS			
<u>TOFACITINIB CITRATE - XELJANZ</u>						
N 213082 001	RE41783	Dec 08, 2025	DS			
<u>TOFERSEN - QALSODY</u>						
N 215887 001	10385341	Apr 01, 2035	DS DP		NCE	Apr 25, 2028
	10669546	Apr 01, 2035	U-3575		ODE-432	Apr 25, 2030
	10968453	Apr 01, 2035	U-3575			
<u>TOLVAPTAN - SAMSCA</u>						
N 022275 001	10905694	Apr 07, 2030	DP			
	8501730	Sep 01, 2026	DS			
<u>TOLVAPTAN - SAMSCA</u>						
N 022275 002	10905694	Apr 07, 2030	DP			
	8501730	Sep 01, 2026	DS			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>TOLVAPTAN - SAMSCA</u>						
N 022275	002	10905694	Apr 07, 2030	DP		
	8501730	Sep 01, 2026	DS			
<u>TOLVAPTAN - SAMSCA</u>						
N 022275	003	10905694	Apr 07, 2030	DP		
	8501730	Sep 01, 2026	DS			
<u>TOLVAPTAN - JYNARQUE</u>						
N 204441	001	10905694	Apr 07, 2030	DP	ODE-178	Apr 23, 2025
	8501730	Sep 01, 2026	DS			
<u>TOLVAPTAN - JYNARQUE</u>						
N 204441	002	10905694	Apr 07, 2030	DP	ODE-178	Apr 23, 2025
	8501730	Sep 01, 2026	DS			
<u>TOLVAPTAN - JYNARQUE</u>						
N 204441	003	10905694	Apr 07, 2030	DP	ODE-178	Apr 23, 2025
	8501730	Sep 01, 2026	DS			
<u>TOLVAPTAN - JYNARQUE</u>						
N 204441	004	10905694	Apr 07, 2030	DP	ODE-178	Apr 23, 2025
	8501730	Sep 01, 2026	DS			
<u>TOLVAPTAN - JYNARQUE</u>						
N 204441	005	10905694	Apr 07, 2030	DP	ODE-178	Apr 23, 2025
	8501730	Sep 01, 2026	DS			
<u>TOPIRAMATE - TROKENDI XR</u>						
N 201635	001	10314790	Nov 16, 2027	DP U-1675		
		10314790	Nov 16, 2027	DP U-1992		
		8298576	Apr 04, 2028	DP U-106		
		8298576	Apr 04, 2028	DP U-1992		
		8298580	Nov 16, 2027	DP U-106		
		8298580	Nov 16, 2027	DP U-1992		
		8663683	Nov 16, 2027	DP U-106		
		8663683	Nov 16, 2027	DP U-1992		
		8877248	Nov 16, 2027	DP U-106		
		8877248	Nov 16, 2027	DP U-1992		
		8889191	Nov 16, 2027	U-106		
		8889191	Nov 16, 2027	U-1992		
		8992989	Nov 16, 2027	DP U-1675		
		8992989	Nov 16, 2027	DP U-1992		
		9549940	Nov 16, 2027	DP U-1675		
		9549940	Nov 16, 2027	DP U-1992		
		9555004	Nov 16, 2027	DP U-1675		
		9555004	Nov 16, 2027	DP U-1992		
		9622983	Nov 16, 2027	DP U-1675		
		9622983	Nov 16, 2027	DP U-1992		
<u>TOPIRAMATE - TROKENDI XR</u>						
N 201635	002	10314790	Nov 16, 2027	DP U-1675		
		10314790	Nov 16, 2027	DP U-1992		
		8298576	Apr 04, 2028	DP U-106		
		8298576	Apr 04, 2028	DP U-1992		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>TOPIRAMATE - TROKENDI XR</u>						
N 201635	002	8298580	Nov 16, 2027	DP U-106		
		8298580	Nov 16, 2027	DP U-1992		
		8663683	Nov 16, 2027	DP U-106		
		8663683	Nov 16, 2027	DP U-1992		
		8877248	Nov 16, 2027	DP U-106		
		8877248	Nov 16, 2027	DP U-1992		
		8889191	Nov 16, 2027	U-106		
		8889191	Nov 16, 2027	U-1992		
		8992989	Nov 16, 2027	DP U-1675		
		8992989	Nov 16, 2027	DP U-1992		
		9549940	Nov 16, 2027	DP U-1675		
		9549940	Nov 16, 2027	DP U-1992		
		9555004	Nov 16, 2027	DP U-1675		
		9555004	Nov 16, 2027	DP U-1992		
		9622983	Nov 16, 2027	DP U-1675		
		9622983	Nov 16, 2027	DP U-1992		
<u>TOPIRAMATE - TROKENDI XR</u>						
N 201635	003	10314790	Nov 16, 2027	DP U-1675		
		10314790	Nov 16, 2027	DP U-1992		
		8298576	Apr 04, 2028	DP U-106		
		8298576	Apr 04, 2028	DP U-1992		
		8298580	Nov 16, 2027	DP U-106		
		8298580	Nov 16, 2027	DP U-1992		
		8663683	Nov 16, 2027	DP U-106		
		8663683	Nov 16, 2027	DP U-1992		
		8877248	Nov 16, 2027	DP U-106		
		8877248	Nov 16, 2027	DP U-1992		
		8889191	Nov 16, 2027	U-106		
		8889191	Nov 16, 2027	U-1992		
		8992989	Nov 16, 2027	DP U-1675		
		8992989	Nov 16, 2027	DP U-1992		
		9549940	Nov 16, 2027	DP U-1675		
		9549940	Nov 16, 2027	DP U-1992		
		9555004	Nov 16, 2027	DP U-1675		
		9555004	Nov 16, 2027	DP U-1992		
		9622983	Nov 16, 2027	DP U-1675		
		9622983	Nov 16, 2027	DP U-1992		
<u>TOPIRAMATE - TROKENDI XR</u>						
N 201635	004	10314790	Nov 16, 2027	DP U-1675		
		10314790	Nov 16, 2027	DP U-1992		
		8298576	Apr 04, 2028	DP U-106		
		8298576	Apr 04, 2028	DP U-1992		
		8298580	Nov 16, 2027	DP U-106		
		8298580	Nov 16, 2027	DP U-1992		
		8663683	Nov 16, 2027	DP U-106		
		8663683	Nov 16, 2027	DP U-1992		
		8877248	Nov 16, 2027	DP U-106		
		8877248	Nov 16, 2027	DP U-1992		
		8889191	Nov 16, 2027	U-106		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>TOPIRAMATE - TROKENDI XR</u>						
N 201635	004 8889191	Nov 16, 2027				U-1992
	8992989	Nov 16, 2027	DP			U-1675
	8992989	Nov 16, 2027	DP			U-1992
	9549940	Nov 16, 2027	DP			U-1675
	9549940	Nov 16, 2027	DP			U-1992
	9555004	Nov 16, 2027	DP			U-1675
	9555004	Nov 16, 2027	DP			U-1992
	9622983	Nov 16, 2027	DP			U-1675
	9622983	Nov 16, 2027	DP			U-1992
<u>TOPIRAMATE - OUDEXY XR</u>						
N 205122	001 10363224	Mar 19, 2033				U-766
	8652527	Mar 19, 2033	DP			
	8889190	Mar 19, 2033	DP			
	9101545	Mar 19, 2033	DP			
	9555005	Mar 19, 2033	DP			
<u>TOPIRAMATE - OUDEXY XR</u>						
N 205122	002 10363224	Mar 19, 2033				U-766
	8652527	Mar 19, 2033	DP			
	8889190	Mar 19, 2033	DP			
	9101545	Mar 19, 2033	DP			
	9555005	Mar 19, 2033	DP			
<u>TOPIRAMATE - OUDEXY XR</u>						
N 205122	003 10363224	Mar 19, 2033				U-766
	8652527	Mar 19, 2033	DP			
	8889190	Mar 19, 2033	DP			
	9101545	Mar 19, 2033	DP			
	9555005	Mar 19, 2033	DP			
<u>TOPIRAMATE - OUDEXY XR</u>						
N 205122	004 10363224	Mar 19, 2033				U-766
	8652527	Mar 19, 2033	DP			
	8889190	Mar 19, 2033	DP			
	9101545	Mar 19, 2033	DP			
	9555005	Mar 19, 2033	DP			
<u>TOPIRAMATE - OUDEXY XR</u>						
N 205122	005 10363224	Mar 19, 2033				U-766
	8652527	Mar 19, 2033	DP			
	8889190	Mar 19, 2033	DP			
	9101545	Mar 19, 2033	DP			
	9555005	Mar 19, 2033	DP			
<u>TOPIRAMATE - EPRONTIA</u>						
N 214679	001 11433046	Aug 21, 2040				U-3413
	11433046	Aug 21, 2040				U-3414
	11433046	Aug 21, 2040				U-3415
	11633374	Aug 21, 2040	DP			
	11826343	Aug 21, 2040	DP			
	11911362	Aug 21, 2040	DP			U-3413
	11911362	Aug 21, 2040	DP			U-3414

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>TOPIRAMATE - EPRONTIA</u>						
N 214679	001 11911362	Aug 21, 2040	DP	U-3415		
<u>TORSEMIDE - SOAANZ</u>						
N 213218	001 10154963	Oct 06, 2033	DP			
<u>TORSEMIDE - SOAANZ</u>						
N 213218	002 10154963	Oct 06, 2033	DP			
<u>TOVORAFENIB - OJEMDA</u>						
N 217700	001 10426782	Jun 23, 2035	DP		NCE	Apr 23, 2029
	8293752	Aug 04, 2031	DS DP		ODE-478	Apr 23, 2031
<u>TOVORAFENIB - OJEMDA</u>						
N 218033	001 10426782	Jun 23, 2035	DP		NCE	Apr 23, 2029
	8293752	Aug 04, 2031	DS DP		ODE-478	Apr 23, 2031
<u>TRABECTEDIN - YONDELIS</u>						
N 207953	001 8895557	Jan 07, 2028	DP			
	8895557*PED	Jul 07, 2028				
<u>TRAMADOL HYDROCHLORIDE - QDOLO</u>						
N 214044	001 11103452	Sep 01, 2040	DP	U-3197		
	11752103	Sep 01, 2040	DP	U-3197		
<u>TRAMETINIB DIMETHYL SULFOXIDE - MEKINIST</u>						
N 204114	001 10869869	Aug 30, 2033		U-3184	I-895	Jun 22, 2025
	10869869*PED	Feb 28, 2034			I-908	Mar 16, 2026
	7378423	May 29, 2027	DS DP		ODE-182	Apr 30, 2025
	7378423*PED	Nov 29, 2027			ODE-183	May 04, 2025
	8580304	Jan 28, 2032	DP		ODE-428	Mar 16, 2030
	8580304*PED	Jul 28, 2032			PED	Oct 30, 2025
	8703781	Oct 15, 2030	DS DP	U-1712	PED	Nov 04, 2025
	8703781	Oct 15, 2030	DS DP	U-2020	PED	Dec 22, 2025
	8703781	Oct 15, 2030	DS DP	U-2037	PED	Sep 16, 2026
	8703781	Oct 15, 2030	DS DP	U-2302	PED	Sep 16, 2030
	8703781	Oct 15, 2030	DS DP	U-2305		
	8703781*PED	Apr 15, 2031				
	8835443	Jun 10, 2025		U-1581		
	8835443	Jun 10, 2025		U-1582		
	8835443	Jun 10, 2025		U-2020		
	8835443	Jun 10, 2025		U-2037		
	8835443	Jun 10, 2025		U-2302		
	8835443	Jun 10, 2025		U-2305		
	8835443*PED	Dec 10, 2025				
	8952018	Oct 15, 2030		U-2020		
	8952018*PED	Apr 15, 2031				
	9155706	Jan 28, 2032	DP			
	9155706*PED	Jul 28, 2032				
	9271941	Jan 28, 2032	DP			
	9271941*PED	Jul 28, 2032				
	9399021	Jan 28, 2032	DP			
	9399021*PED	Jul 28, 2032				

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>TRAMETINIB DIMETHYL SULFOXIDE - MEKINIST</u>						
N 204114 002	10869869	Aug 30, 2033	U-3184		I-895	Jun 22, 2025
	10869869*PED	Feb 28, 2034			I-908	Mar 16, 2026
	7378423	May 29, 2027	DS DP		ODE-182	Apr 30, 2025
	7378423*PED	Nov 29, 2027			ODE-183	May 04, 2025
	8580304	Jan 28, 2032	DP		ODE-428	Mar 16, 2030
	8580304*PED	Jul 28, 2032			PED	Oct 30, 2025
	8703781	Oct 15, 2030	DS DP U-1712		PED	Nov 04, 2025
	8703781	Oct 15, 2030	DS DP U-2020		PED	Dec 22, 2025
	8703781	Oct 15, 2030	DS DP U-2037		PED	Sep 16, 2026
	8703781	Oct 15, 2030	DS DP U-2302		PED	Sep 16, 2030
	8703781	Oct 15, 2030	DS DP U-2305			
	8703781*PED	Apr 15, 2031				
	8835443	Jun 10, 2025	U-1581			
	8835443	Jun 10, 2025	U-1582			
	8835443	Jun 10, 2025	U-2020			
	8835443	Jun 10, 2025	U-2037			
	8835443	Jun 10, 2025	U-2302			
	8835443	Jun 10, 2025	U-2305			
	8835443*PED	Dec 10, 2025				
	8952018	Oct 15, 2030	U-2020			
	8952018*PED	Apr 15, 2031				
	9155706	Jan 28, 2032	DP			
	9155706*PED	Jul 28, 2032				
	9271941	Jan 28, 2032	DP			
	9271941*PED	Jul 28, 2032				
	9399021	Jan 28, 2032	DP			
	9399021*PED	Jul 28, 2032				
<u>TRAMETINIB DIMETHYL SULFOXIDE - MEKINIST</u>						
N 204114 003	10869869	Aug 30, 2033	U-3184		I-895	Jun 22, 2025
	10869869*PED	Feb 28, 2034			I-908	Mar 16, 2026
	7378423	May 29, 2027	DS DP		ODE-182	Apr 30, 2025
	7378423*PED	Nov 29, 2027			ODE-183	May 04, 2025
	8580304	Jan 28, 2032	DP		ODE-428	Mar 16, 2030
	8580304*PED	Jul 28, 2032			PED	Oct 30, 2025
	8703781	Oct 15, 2030	DS DP U-1712		PED	Nov 04, 2025
	8703781	Oct 15, 2030	DS DP U-2020		PED	Dec 22, 2025
	8703781	Oct 15, 2030	DS DP U-2037		PED	Sep 16, 2026
	8703781	Oct 15, 2030	DS DP U-2302		PED	Sep 16, 2030
	8703781	Oct 15, 2030	DS DP U-2305			
	8703781*PED	Apr 15, 2031				
	8835443	Jun 10, 2025	U-1581			
	8835443	Jun 10, 2025	U-1582			
	8835443	Jun 10, 2025	U-2020			
	8835443	Jun 10, 2025	U-2037			
	8835443	Jun 10, 2025	U-2302			
	8835443	Jun 10, 2025	U-2305			
	8835443*PED	Dec 10, 2025				
	8952018	Oct 15, 2030	U-2020			
	8952018*PED	Apr 15, 2031				

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>TRAMETINIB DIMETHYL SULFOXIDE - MEKINIST</u>						
N 204114	003 9155706	Jan 28, 2032	DP			
	9155706*PED	Jul 28, 2032				
	9271941	Jan 28, 2032	DP			
	9271941*PED	Jul 28, 2032				
	9399021	Jan 28, 2032	DP			
	9399021*PED	Jul 28, 2032				
<u>TRAMETINIB DIMETHYL SULFOXIDE - MEKINIST</u>						
N 217513	001 7378423	May 29, 2027	DS DP		NP	Mar 16, 2026
	7378423*PED	Nov 29, 2027			ODE-428	Mar 16, 2030
	8703781	Oct 15, 2030	DS DP U-3564		PED	Sep 16, 2026
	8703781*PED	Apr 15, 2031			PED	Sep 16, 2030
	8835443	Jun 10, 2025	U-3564			
	8835443*PED	Dec 10, 2025				
<u>TRANEXAMIC ACID - LYSTEDA</u>						
N 022430	001 7947739	Mar 04, 2025	DP			
	8022106	Mar 04, 2025	U-1182			
	8273795	Mar 04, 2025	U-1182			
	8487005	Mar 04, 2025	DP U-1182			
	8791160	Mar 04, 2025	DP U-1182			
	8809394	Mar 04, 2025	DP U-1182			
	8957113	Mar 04, 2025	DP U-1182			
	9060939	Mar 04, 2025	DP			
<u>TRAVOPROST - TRAVATAN Z</u>						
N 021994	001 8268299	Oct 13, 2029	DP			
	8323630	Sep 20, 2027	DP			
	8388941	Sep 20, 2027	DP			
<u>TRAVOPROST - IZBA</u>						
N 204822	001 8178582	Oct 10, 2029	DP			
	8722735	Oct 10, 2029	DP			
	8754123	May 19, 2029	DP			
	9144561	Mar 13, 2029	DP			
<u>TRAVOPROST - IDOSE TR</u>						
N 218010	001 10206813	Oct 17, 2030	DP		NP	Dec 13, 2026
	11426306	Oct 17, 2030	DP			
<u>TRAZODONE HYDROCHLORIDE - DESYREL</u>						
N 018207	001 8133893	Mar 13, 2029	DS DP			
<u>TRAZODONE HYDROCHLORIDE - DESYREL</u>						
N 018207	002 8133893	Mar 13, 2029	DS DP			
<u>TRAZODONE HYDROCHLORIDE - DESYREL</u>						
N 018207	003 8133893	Mar 13, 2029	DS DP			
<u>TRAZODONE HYDROCHLORIDE - DESYREL</u>						
N 018207	004 8133893	Mar 13, 2029	DS DP			
<u>TRAZODONE HYDROCHLORIDE - OLEPTRO</u>						
N 022411	001 7829120	Mar 27, 2027	DP U-796			
	8133893	Mar 13, 2029	DS DP			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>TRAZODONE HYDROCHLORIDE - OLEPTRO</u>						
N 022411	002 7829120	Mar 27, 2027	DP U-796			
	8133893	Mar 13, 2029	DS DP			
<u>TRAZODONE HYDROCHLORIDE - RALDESY</u>						
N 218637	001 8133893	Mar 13, 2029	DS DP			
<u>TREPROSTINIL - REMODULIN</u>						
N 021272	001 11723887	Dec 15, 2028	DS			
	7999007	Mar 29, 2029	DP U-1437			
	8653137	Sep 05, 2028	U-1437			
	8658694	Sep 05, 2028	U-1437			
	9593066	Dec 15, 2028	DS			
<u>TREPROSTINIL - REMODULIN</u>						
N 021272	002 11723887	Dec 15, 2028	DS			
	7999007	Mar 29, 2029	DP U-1437			
	8653137	Sep 05, 2028	U-1437			
	8658694	Sep 05, 2028	U-1437			
	9593066	Dec 15, 2028	DS			
<u>TREPROSTINIL - REMODULIN</u>						
N 021272	003 11723887	Dec 15, 2028	DS			
	7999007	Mar 29, 2029	DP U-1437			
	8653137	Sep 05, 2028	U-1437			
	8658694	Sep 05, 2028	U-1437			
	9593066	Dec 15, 2028	DS			
<u>TREPROSTINIL - REMODULIN</u>						
N 021272	004 11723887	Dec 15, 2028	DS			
	7999007	Mar 29, 2029	DP U-1437			
	8653137	Sep 05, 2028	U-1437			
	8658694	Sep 05, 2028	U-1437			
	9593066	Dec 15, 2028	DS			
<u>TREPROSTINIL - REMODULIN</u>						
N 021272	005 11723887	Dec 15, 2028	DS			
	7999007	Mar 29, 2029	DP U-1437			
	8653137	Sep 05, 2028	U-1437			
	8658694	Sep 05, 2028	U-1437			
	9593066	Dec 15, 2028	DS			
<u>TREPROSTINIL - REMODULIN</u>						
N 021272	006 11723887	Dec 15, 2028	DS			
	7999007	Mar 29, 2029	DP U-1437			
	8653137	Sep 05, 2028	U-1437			
	8658694	Sep 05, 2028	U-1437			
	9593066	Dec 15, 2028	DS			
<u>TREPROSTINIL - REMODULIN</u>						
N 021272	007 11723887	Dec 15, 2028	DS			
	7999007	Mar 29, 2029	DP U-1437			
	8653137	Sep 05, 2028	U-1437			
	8658694	Sep 05, 2028	U-1437			
	9593066	Dec 15, 2028	DS			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>TREPROSTINIL - REMODULIN</u>						
N 021272	008	11723887	Dec 15, 2028	DS		
		7999007	Mar 29, 2029	DP	U-1437	
		8653137	Sep 05, 2028		U-1437	
		8658694	Sep 05, 2028		U-1437	
		9593066	Dec 15, 2028	DS		
<u>TREPROSTINIL - TYVASO</u>						
N 022387	001	10376525	May 14, 2027		U-1849	
		11723887	Dec 15, 2028	DS		
		11826327	Feb 03, 2042		U-3749	
		9339507	Mar 10, 2028	DP		
		9358240	May 05, 2028		U-1849	
		9593066	Dec 15, 2028	DS		
<u>TREPROSTINIL - REMODULIN</u>						
N 208276	001	9593066	Dec 15, 2028	DS		
<u>TREPROSTINIL - REMODULIN</u>						
N 208276	002	9593066	Dec 15, 2028	DS		
<u>TREPROSTINIL - REMODULIN</u>						
N 208276	003	9593066	Dec 15, 2028	DS		
<u>TREPROSTINIL - REMODULIN</u>						
N 208276	004	9593066	Dec 15, 2028	DS		
<u>TREPROSTINIL - TYVASO DPI</u>						
N 214324	001	10130685	Aug 23, 2025	DP	NP	May 23, 2025
		10421729	Apr 01, 2035	DP		
		10772883	Jun 11, 2030	DP		
		11723887	Dec 15, 2028	DS		
		11826327	Feb 03, 2042		U-3749	
		9593066	Dec 15, 2028	DS		
<u>TREPROSTINIL - TYVASO DPI</u>						
N 214324	002	10130685	Aug 23, 2025	DP	NP	May 23, 2025
		10421729	Apr 01, 2035	DP		
		10772883	Jun 11, 2030	DP		
		11723887	Dec 15, 2028	DS		
		11826327	Feb 03, 2042		U-3749	
		9593066	Dec 15, 2028	DS		
<u>TREPROSTINIL - TYVASO DPI</u>						
N 214324	003	10130685	Aug 23, 2025	DP	NP	May 23, 2025
		10421729	Apr 01, 2035	DP		
		10772883	Jun 11, 2030	DP		
		11723887	Dec 15, 2028	DS		
		11826327	Feb 03, 2042		U-3749	
		9593066	Dec 15, 2028	DS		
<u>TREPROSTINIL - TYVASO DPI</u>						
N 214324	004	10130685	Aug 23, 2025	DP	NP	May 23, 2025
		10421729	Apr 01, 2035	DP		
		10772883	Jun 11, 2030	DP		
		11723887	Dec 15, 2028	DS		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>TREPROSTINIL - TYVASO DPI</u>						
N 214324 004	11826327	Feb 03, 2042	U-3749			
	9593066	Dec 15, 2028	DS			
<u>TREPROSTINIL - TYVASO DPI</u>						
N 214324 005	10130685	Aug 23, 2025	DP			
	10421729	Apr 01, 2035	DP			
	10772883	Jun 11, 2030	DP			
	11723887	Dec 15, 2028	DS			
	11826327	Feb 03, 2042	U-3749			
	9593066	Dec 15, 2028	DS			
<u>TREPROSTINIL DIOLAMINE - ORENITRAM</u>						
N 203496 001	11723887	Dec 15, 2028	DS		ODE-272	Oct 18, 2026
	7417070	Jul 30, 2026	DS			
	8349892	Jan 22, 2031	DP			
	8410169	Feb 13, 2030	DP			
	8747897	Aug 11, 2031	DP U-2724			
	8747897	Aug 11, 2031	DP U-2725			
	9393203	Apr 27, 2026	DP U-1877			
	9593066	Dec 15, 2028	DS			
	9604901	Dec 15, 2028	DS	Y		
<u>TREPROSTINIL DIOLAMINE - ORENITRAM</u>						
N 203496 002	11723887	Dec 15, 2028	DS		ODE-272	Oct 18, 2026
	7417070	Jul 30, 2026	DS			
	8349892	Jan 22, 2031	DP			
	8410169	Feb 13, 2030	DP			
	8497393	Dec 15, 2028	DS	Y		
	8747897	Aug 11, 2031	DP U-2724			
	8747897	Aug 11, 2031	DP U-2725			
	9393203	Apr 27, 2026	DP U-1877			
	9593066	Dec 15, 2028	DS			
<u>TREPROSTINIL DIOLAMINE - ORENITRAM</u>						
N 203496 003	11723887	Dec 15, 2028	DS		ODE-272	Oct 18, 2026
	7417070	Jul 30, 2026	DS			
	8349892	Jan 22, 2031	DP			
	8410169	Feb 13, 2030	DP			
	8497393	Dec 15, 2028	DS	Y		
	8747897	Aug 11, 2031	DP U-2724			
	8747897	Aug 11, 2031	DP U-2725			
	9393203	Apr 27, 2026	DP U-1877			
	9593066	Dec 15, 2028	DS			
<u>TREPROSTINIL DIOLAMINE - ORENITRAM</u>						
N 203496 004	11723887	Dec 15, 2028	DS		ODE-272	Oct 18, 2026
	7417070	Jul 30, 2026	DS			
	8349892	Jan 22, 2031	DP			
	8410169	Feb 13, 2030	DP			
	8497393	Dec 15, 2028	DS	Y		
	8747897	Aug 11, 2031	DP U-2724			
	8747897	Aug 11, 2031	DP U-2725			
	9393203	Apr 27, 2026	DP U-1877			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>TREPROSTINIL DIOLAMINE - ORENITRAM</u>						
N 203496	004	9593066	Dec 15, 2028	DS		
<u>TREPROSTINIL DIOLAMINE - ORENITRAM</u>						
N 203496	005	11723887	Dec 15, 2028	DS	ODE-272	Oct 18, 2026
		7417070	Jul 30, 2026	DS		
		8349892	Jan 22, 2031	DP		
		8410169	Feb 13, 2030	DP		
		8747897	Aug 11, 2031	DP U-2724		
		8747897	Aug 11, 2031	DP U-2725		
		9393203	Apr 27, 2026	DP U-1877		
		9593066	Dec 15, 2028	DS		
		9604901	Dec 15, 2028	DS	Y	
<u>TRETINOIN - ALTRENO</u>						
N 209353	001	10653656	Aug 22, 2038	DP U-2368		
		11324710	Aug 22, 2038	U-2368		
<u>TRIAMCINOLONE ACETONIDE - TRIESENC</u>						
N 022048	001	8128960	Dec 17, 2029	DP		
		8211880	Mar 10, 2029	U-1257		
		8211880	Mar 10, 2029	U-1258		
<u>TRIAMCINOLONE ACETONIDE - ZILRETTA</u>						
N 208845	001	8828440	Aug 04, 2031	DP		
		9555048	Aug 04, 2031	U-2151		
<u>TRIAMCINOLONE ACETONIDE - XIPERE</u>						
N 211950	001	8636713	May 02, 2027	U-3234		
		9636332	Nov 08, 2033	U-3234		
		9937075	May 02, 2034	DP		
<u>TRICLABENDAZOLE - EGATEN</u>						
N 208711	001				ODE-228	Feb 13, 2026
<u>TRIENTINE TETRAHYDROCHLORIDE - CUVRIOR</u>						
N 215760	001	10988436	May 03, 2039	DS	NP	Apr 28, 2025
		11072577	May 03, 2039	U-3370	ODE-401	Apr 28, 2029
<u>TRIFAROTENE - AKLIEF</u>						
N 211527	001	7807708	Jul 19, 2031	DS DP		
		8227507	Dec 21, 2025	U-818		
		8470871	Dec 21, 2025	U-2639		
		9084778	May 30, 2033	DP U-134		
		9498465	May 30, 2033	DP U-1033		
<u>TRIEPTANOIN - DOJOLVI</u>						
N 213687	001	8697748	Apr 28, 2029	DP	NCE	Jun 30, 2025
		9186344	Jul 01, 2025	DP	ODE-311	Jun 30, 2027
<u>TRILACICLIB DIHYDROCHLORIDE - COSELA</u>						
N 214200	001	10085992	Mar 14, 2034	U-3081	NCE	Feb 12, 2026
		10189849	Oct 25, 2031	DS		
		10189850	Oct 25, 2031	DP		
		10927120	Oct 25, 2031	DP		
		10966984	Mar 14, 2034	U-3079		
		10966984	Mar 14, 2034	U-3080		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>TRILACICLIB DIHYDROCHLORIDE - COSELA</u>						
N 214200	001	11040042	Oct 25, 2031	DP		
		11529352	Jul 23, 2039	U-3504		
		11717523	Mar 14, 2034	U-3678		
		11717523	Mar 14, 2034	U-3679		
		8598186	Oct 25, 2031	DS DP		
		8598197	Oct 25, 2031	DS DP		
		9487530	Mar 14, 2034	U-3079		
		9487530	Mar 14, 2034	U-3080		
		9957276	Oct 25, 2031	DS		
<u>TRIPTORELIN PAMOATE - TRELSTAR</u>						
N 022437	001	10166181	Jun 30, 2029	DP		
<u>TRIPTORELIN PAMOATE - TRIPTODUR KIT</u>						
N 208956	001	10166181	Jun 30, 2029	DP		
<u>TROFINETIDE - DAYBUE</u>						
N 217026	001	11370755	Aug 03, 2040	DS DP	NCE	Mar 10, 2028
		11827600	Jul 12, 2042	DS DP U-3556	ODE-425	Mar 10, 2030
		9212204	Jan 27, 2032	U-3556		
<u>TROMETHAMINE - TROMETHAMINE</u>						
A 213116	001				CGT	Jul 01, 2025
<u>TROSPIUM CHLORIDE; XANOMELINE TARTRATE - COBENFY</u>						
N 216158	001	10238643	Jul 21, 2030	DP	NCE	Sep 26, 2029
		10265311	Jul 21, 2030	U-3513		
		10369143	Jul 21, 2030	U-3513		
		10369144	Jul 21, 2030	DP U-3513		
		10695339	Jul 21, 2030	U-3513		
		10925832	Sep 27, 2039	DP		
		10933020	Sep 27, 2039	DP U-3513		
		11452692	Sep 27, 2039	U-3513		
		11471413	Sep 27, 2039	U-3513		
		11890378	Sep 27, 2039	DP U-3513		
<u>TROSPIUM CHLORIDE; XANOMELINE TARTRATE - COBENFY</u>						
N 216158	002	10238643	Jul 21, 2030	DP	NCE	Sep 26, 2029
		10265311	Jul 21, 2030	U-3513		
		10369143	Jul 21, 2030	U-3513		
		10369144	Jul 21, 2030	DP U-3513		
		10695339	Jul 21, 2030	U-3513		
		10925832	Sep 27, 2039	DP		
		10933020	Sep 27, 2039	DP U-3513		
		11452692	Sep 27, 2039	U-3513		
		11471413	Sep 27, 2039	U-3513		
		11890378	Sep 27, 2039	DP U-3513		
<u>TROSPIUM CHLORIDE; XANOMELINE TARTRATE - COBENFY</u>						
N 216158	003	10238643	Jul 21, 2030	DP	NCE	Sep 26, 2029
		10265311	Jul 21, 2030	U-3513		
		10369143	Jul 21, 2030	U-3513		
		10369144	Jul 21, 2030	DP U-3513		
		10695339	Jul 21, 2030	U-3513		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>TROSPIUM CHLORIDE; XANOMELINE TARTRATE - COBENFY</u>						
N 216158	003	10925832	Sep 27, 2039	DP		
		10933020	Sep 27, 2039	DP U-3513		
		11452692	Sep 27, 2039	U-3513		
		11471413	Sep 27, 2039	U-3513		
		11890378	Sep 27, 2039	DP U-3513		
<u>TUCATINIB - TUKYSA</u>						
N 213411	001	11207324	Apr 27, 2038	U-3783	I-906	Jan 19, 2026
		11504370	Mar 25, 2033	U-2788	NCE	Apr 17, 2025
		11666572	Apr 27, 2038	U-3783	ODE-309	Apr 17, 2027
		12048698	Apr 27, 2038	U-3783	ODE-422	Jan 19, 2030
		8648087	Apr 12, 2031	DS DP		
		9457093	Oct 12, 2032	DP U-2788		
		9693989	May 09, 2027	DP U-2788		
<u>TUCATINIB - TUKYSA</u>						
N 213411	002	11207324	Apr 27, 2038	U-3783	I-906	Jan 19, 2026
		11504370	Mar 25, 2033	U-2788	NCE	Apr 17, 2025
		11666572	Apr 27, 2038	U-3783	ODE-309	Apr 17, 2027
		12048698	Apr 27, 2038	U-3783	ODE-422	Jan 19, 2030
		8648087	Apr 12, 2031	DS DP		
		9457093	Oct 12, 2032	DP U-2788		
		9693989	May 09, 2027	DP U-2788		
<u>UBROGEPANT - UBRELVY</u>						
N 211765	001	10117836	Jan 30, 2035	DP		
		11717515	Dec 22, 2041	U-3677		
		11857542	Dec 22, 2041	U-3786		
		11925709	Jan 30, 2035	DP		
		8754096	Jul 19, 2032	DS DP U-2717		
		8912210	Dec 23, 2033	DS DP		
		9499545	Nov 10, 2031	DS DP U-2718		
		9833448	Nov 10, 2031	U-2718		
<u>UBROGEPANT - UBRELVY</u>						
N 211765	002	10117836	Jan 30, 2035	DP		
		11925709	Jan 30, 2035	DP		
		12070450	Dec 22, 2041	U-3992		
		8754096	Jul 19, 2032	DS DP U-2717		
		8912210	Dec 23, 2033	DS DP		
		9499545	Nov 10, 2031	DS DP U-2718		
		9833448	Nov 10, 2031	U-2718		
<u>ULIPRISTAL ACETATE - ELLA</u>						
N 022474	001	10159681	Apr 13, 2030	U-2510		
		10772897	Apr 13, 2030	U-2958		
		8426392	Jun 12, 2030	U-1389		
		8512745	Jun 02, 2030	DP		
		8735380	Feb 20, 2029	DP		
		8962603	Jun 12, 2030	U-1657		
		9283233	Apr 13, 2030	U-1821		
		9844510	Dec 08, 2028	DP		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>UMBRALISIB TOSYLATE - UKONIO</u>						
N 213176 001	10072013	Jul 02, 2033	U-3063		NCE	Feb 05, 2026
	10072013	Jul 02, 2033	U-3064			
	10414773	May 26, 2035	DS DP U-3063			
	10414773	May 26, 2035	DS DP U-3064			
	10570142	Jul 02, 2033	DS DP U-3063			
	10570142	Jul 02, 2033	DS DP U-3064			
	10947244	May 26, 2035	U-3063			
	10947244	May 26, 2035	U-3064			
	10981919	Jul 02, 2033	U-3063			
	10981919	Jul 02, 2033	U-3064			
	9150579	Jul 02, 2033	DS DP			
	9669033	Jul 02, 2033	U-3063			
	9669033	Jul 02, 2033	U-3064			
	9969740	May 26, 2035	DS DP U-3063			
	9969740	May 26, 2035	DS DP U-3064			
<u>UMECLIDINIUM BROMIDE - INCRUSE ELLIPTA</u>						
N 205382 001	7488827	Dec 18, 2027	DS DP			
	7498440	Apr 27, 2025	DS DP			
	8183257	Jul 27, 2025	U-1476			
	8201556	Feb 05, 2029	DP			
	8309572	Apr 27, 2025	U-1476			
	8746242	Oct 11, 2030	DP			
	9333310	Oct 02, 2027	DP			
<u>UMECLIDINIUM BROMIDE; VILANTEROL TRIFENATATE - ANORO ELLIPTA</u>						
N 203975 001	11090294	Nov 29, 2030	U-3203			
	7439393	May 21, 2025	DS DP U-1476			
	7439393*PED	Nov 21, 2025				
	7488827	Dec 18, 2027	DS DP			
	7498440	Apr 27, 2025	DS DP			
	8161968	Feb 05, 2028	DP			
	8161968*PED	Aug 05, 2028				
	8183257	Jul 27, 2025	U-1476			
	8309572	Apr 27, 2025	U-1476			
	8511304	Jun 14, 2027	DP U-1476			
	8511304*PED	Dec 14, 2027				
	8534281	Mar 08, 2030	DP			
	8534281*PED	Sep 08, 2030				
	8746242	Oct 11, 2030	DP			
	8746242*PED	Apr 11, 2031				
	9333310	Oct 02, 2027	DP			
	9333310*PED	Apr 02, 2028				
	9750726	Nov 29, 2030	DP			
<u>UPADACITINIB - RINVOO</u>						
N 211675 001	10519164	Oct 17, 2036	DP		I-883	Jan 14, 2025
	10597400	Oct 17, 2036	U-3255		I-886	Mar 16, 2025
	10981923	Oct 17, 2036	DS		I-888	Apr 29, 2025
	10981924	Oct 17, 2036	DP		I-919	May 18, 2026
	10995095	Oct 17, 2036	U-3298		I-946	Apr 26, 2027

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>UPADACITINIB - RINVOO</u>						
N 211675 001	11186584	Oct 17, 2036	DS		NPP	Apr 26, 2027
	11198697	Oct 17, 2036	DP		ODE-481	Apr 26, 2031
	11365198	Oct 17, 2036		U-3275		
	11365198	Oct 17, 2036		U-3946		
	11512092	Oct 17, 2036		U-3275		
	11512092	Oct 17, 2036		U-3371		
	11512092	Oct 17, 2036		U-3487		
	11512092	Oct 17, 2036		U-3946		
	11524964	Oct 17, 2036		U-3371		
	11535624	Oct 17, 2036		U-3255		
	11535625	Oct 17, 2036		U-3298		
	11564922	Mar 09, 2038		U-3624		
	11607411	Mar 09, 2038		U-3341		
	11661425	Oct 17, 2036	DS DP			
	11680069	Oct 17, 2036	DS DP			
	11718627	Oct 17, 2036	DS DP			
	11767326	Oct 17, 2036		U-3298		
	11773105	Oct 17, 2036	DS DP			
	11773106	Oct 17, 2036		U-3275		
	11773106	Oct 17, 2036		U-3371		
	11773106	Oct 17, 2036		U-3487		
	11773106	Oct 17, 2036		U-3946		
	11780847	Oct 17, 2036	DP			
	11780848	Oct 17, 2036		U-3371		
	11787815	Oct 17, 2036	DP			
	11795175	Oct 17, 2036		U-3255		
	11976077	Oct 17, 2036		U-3298		
	11993605	Oct 17, 2036		U-3275		
	11993605	Oct 17, 2036		U-3946		
	11993606	Oct 17, 2036		U-3487		
	12077545	Oct 17, 2036	DP			
	12103933	Oct 17, 2036	DP			
	12110297	Oct 17, 2036	DP			
	12110298	Oct 17, 2036	DS DP			
	8962629	Jan 15, 2031	DS	U-3255		
	8962629	Jan 15, 2031	DS	U-3275		
	8962629	Jan 15, 2031	DS	U-3341		
	8962629	Jan 15, 2031	DS	U-3371		
	8962629	Jan 15, 2031	DS	U-3624		
	8962629	Jan 15, 2031	DS	U-3945		
	8962629	Jan 15, 2031	DS	U-3946		
	9951080	Oct 17, 2036	DS DP			
	9963459	Oct 17, 2036	DP			
	RE47221	Aug 16, 2033	DS			
<u>UPADACITINIB - RINVOO</u>						
N 211675 002	10344036	Oct 17, 2036	DP		I-883	Jan 14, 2025
	10550126	Oct 17, 2036		U-3298	I-886	Mar 16, 2025
	10730883	Oct 17, 2036	DP		I-919	May 18, 2026
	10981923	Oct 17, 2036	DS		I-946	Apr 26, 2027

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>UPADACITINIB - RINVOO</u>						
N 211675 002	10981924	Oct 17, 2036	DP		NPP	Apr 26, 2027
	11186584	Oct 17, 2036	DS		ODE-481	Apr 26, 2031
	11198697	Oct 17, 2036	DP			
	11535626	Oct 17, 2036		U-3298		
	11564922	Mar 09, 2038		U-3624		
	11607411	Mar 09, 2038		U-3341		
	11661425	Oct 17, 2036	DS DP			
	11680069	Oct 17, 2036	DS DP			
	11718627	Oct 17, 2036	DS DP			
	11767326	Oct 17, 2036		U-3298		
	11773105	Oct 17, 2036	DS DP			
	12077545	Oct 17, 2036	DP			
	12091415	Oct 17, 2036	DP			
	12110298	Oct 17, 2036	DS DP			
	12116373	Oct 17, 2036	DP			
	8962629	Jan 15, 2031	DS	U-3341		
	8962629	Jan 15, 2031	DS	U-3624		
	9951080	Oct 17, 2036	DS DP			
	RE47221	Aug 16, 2033	DS			
<u>UPADACITINIB - RINVOO</u>						
N 211675 003	10202393	Oct 17, 2036	DP		I-886	Mar 16, 2025
	10981923	Oct 17, 2036	DS		I-919	May 18, 2026
	11186584	Oct 17, 2036	DS		I-946	Apr 26, 2027
	11198697	Oct 17, 2036	DP		NPP	Apr 26, 2027
	11564922	Mar 09, 2038		U-3624	ODE-481	Apr 26, 2031
	11607411	Mar 09, 2038		U-3341		
	11661425	Oct 17, 2036	DS DP			
	11680069	Oct 17, 2036	DS DP			
	11718627	Oct 17, 2036	DS DP			
	11773105	Oct 17, 2036	DS DP			
	12077545	Oct 17, 2036	DP			
	12110298	Oct 17, 2036	DS DP			
	12134621	Oct 17, 2036	DP			
	8962629	Jan 15, 2031	DS	U-3341		
	8962629	Jan 15, 2031	DS	U-3624		
	9951080	Oct 17, 2036	DS DP			
	RE47221	Aug 16, 2033	DS			
<u>UPADACITINIB - RINVOO LO</u>						
N 218347 001	10981923	Oct 17, 2036	DS		NP	Apr 26, 2027
	11186584	Oct 17, 2036	DS		ODE-481	Apr 26, 2031
	11661425	Oct 17, 2036	DS			
	11680069	Oct 17, 2036	DS			
	11718627	Oct 17, 2036	DS			
	11773105	Oct 17, 2036	DS			
	12110298	Oct 17, 2036	DS			
	8962629	Jan 15, 2031	DS	U-3945		
	8962629	Jan 15, 2031	DS	U-3946		
	9951080	Oct 17, 2036	DS			
	RE47221	Aug 16, 2033	DS			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>UPADACITINIB - RINVOQ LO</u>						
N 218347	001	10981923	Oct 17, 2036	DS	NP	Apr 26, 2027
		11186584	Oct 17, 2036	DS	ODE-481	Apr 26, 2031
		11661425	Oct 17, 2036	DS		
		11680069	Oct 17, 2036	DS		
		11718627	Oct 17, 2036	DS		
		11773105	Oct 17, 2036	DS		
		12110298	Oct 17, 2036	DS		
		8962629	Jan 15, 2031	DS	U-3945	
		8962629	Jan 15, 2031	DS	U-3946	
		9951080	Oct 17, 2036	DS		
		RE47221	Aug 16, 2033	DS		
<u>URIDINE TRIACETATE - VISTOGARD</u>						
N 208159	001	7776838	Aug 17, 2027		U-1791	
<u>VADADUSTAT - VAFSEO</u>						
N 215192	001	10149842	Nov 14, 2034	DS DP	NCE	Mar 27, 2029
		11065237	Nov 14, 2034	DS DP	U-3876	
		11324734	Mar 31, 2036	DP		
		11844756	Mar 31, 2036		U-3876	
		11857543	Jun 09, 2034		U-3876	
		7811595	Mar 13, 2028	DS DP		
		8323671	Apr 03, 2028		U-3876	
		8343952	Aug 14, 2027	DS DP		
		8598210	Jun 26, 2027	DS DP		
		8940773	Jun 26, 2027		U-3876	
		9701636	Nov 14, 2034	DS DP		
		9987262	Nov 14, 2034		U-3876	
		RE47437	Jun 26, 2027	DS DP		
<u>VADADUSTAT - VAFSEO</u>						
N 215192	002	10149842	Nov 14, 2034	DS DP	NCE	Mar 27, 2029
		11065237	Nov 14, 2034	DS DP	U-3876	
		11324734	Mar 31, 2036	DP		
		11844756	Mar 31, 2036		U-3876	
		11857543	Jun 09, 2034		U-3876	
		7811595	Mar 13, 2028	DS DP		
		8323671	Apr 03, 2028		U-3876	
		8343952	Aug 14, 2027	DS DP		
		8598210	Jun 26, 2027	DS DP		
		8940773	Jun 26, 2027		U-3876	
		9701636	Nov 14, 2034	DS DP		
		9987262	Nov 14, 2034		U-3876	
		RE47437	Jun 26, 2027	DS DP		
<u>VADADUSTAT - VAFSEO</u>						
N 215192	003	10149842	Nov 14, 2034	DS DP	NCE	Mar 27, 2029
		11065237	Nov 14, 2034	DS DP	U-3876	
		11324734	Mar 31, 2036	DP		
		11844756	Mar 31, 2036		U-3876	
		11857543	Jun 09, 2034		U-3876	
		7811595	Mar 13, 2028	DS DP		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>VADADUSTAT - VAFSEO</u>						
N 215192	003 8323671	Apr 03, 2028	U-3876			
	8343952	Aug 14, 2027	DS DP			
	8598210	Jun 26, 2027	DS DP			
	8940773	Jun 26, 2027	U-3876			
	9701636	Nov 14, 2034	DS DP			
	9987262	Nov 14, 2034	U-3876			
	RE47437	Jun 26, 2027	DS DP			
<u>VALBENZAZINE TOSYLATE - INGREZZA</u>						
N 209241	001 10065952	Oct 28, 2036	DS DP U-1995		I-925	Aug 18, 2026
	10065952	Oct 28, 2036	DS DP U-3055		ODE-440	Aug 18, 2030
	10844058	Oct 28, 2036	DS DP U-1995			
	10844058	Oct 28, 2036	DS DP U-3055			
	10851103	Oct 28, 2036	DS DP U-1995			
	10851103	Oct 28, 2036	DS DP U-3055			
	10851104	Oct 28, 2036	DS U-1995			
	10851104	Oct 28, 2036	DS U-3055			
	10857137	Oct 10, 2037	U-1995			
	10857148	Oct 10, 2037	U-1995			
	10874648	Oct 10, 2037	U-1995			
	10874648	Oct 10, 2037	U-3046			
	10874648	Oct 10, 2037	U-3055			
	10906902	Dec 22, 2036	DS DP			
	10906903	Dec 22, 2036	DS DP			
	10912771	Oct 10, 2037	U-1995			
	10912771	Oct 10, 2037	U-3055			
	10912771	Oct 10, 2037	U-3076			
	10919892	Dec 22, 2036	DS DP			
	10940141	Aug 10, 2040	U-1995			
	10952997	Oct 10, 2037	U-1995			
	10993941	Oct 10, 2037	U-1995			
	11026931	Aug 14, 2039	U-1995			
	11026939	Sep 18, 2038	DP U-1995			
	11026939	Sep 18, 2038	DP U-3055			
	11040029	Oct 10, 2037	U-1995			
	11311532	Sep 18, 2038	DP U-1995			
	11311532	Sep 18, 2038	DP U-3055			
	11439629	Oct 10, 2037	U-3055			
	11654142	Nov 14, 2038	U-3055			
	8039627	Apr 11, 2031	DS DP			
	8357697	Nov 08, 2027	U-1995			
	8357697	Nov 08, 2027	U-3055			
<u>VALBENZAZINE TOSYLATE - INGREZZA</u>						
N 209241	002 10065952	Oct 28, 2036	DS DP U-1995		I-925	Aug 18, 2026
	10065952	Oct 28, 2036	DS DP U-3055		ODE-440	Aug 18, 2030
	10844058	Oct 28, 2036	DS DP U-1995			
	10844058	Oct 28, 2036	DS DP U-3055			
	10851103	Oct 28, 2036	DS DP U-1995			
	10851103	Oct 28, 2036	DS DP U-3055			
	10851104	Oct 28, 2036	DS U-1995			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>VALBENAZINE TOSYLATE - INGREZZA</u>						
N 209241 002	10851104	Oct 28, 2036	DS U-3055			
	10857137	Oct 10, 2037	U-1995			
	10857148	Oct 10, 2037	U-1995			
	10874648	Oct 10, 2037	U-1995			
	10874648	Oct 10, 2037	U-3046			
	10874648	Oct 10, 2037	U-3055			
	10906902	Dec 22, 2036	DS DP			
	10906903	Dec 22, 2036	DS DP			
	10912771	Oct 10, 2037	U-1995			
	10912771	Oct 10, 2037	U-3055			
	10912771	Oct 10, 2037	U-3076			
	10919892	Dec 22, 2036	DS DP			
	10940141	Aug 10, 2040	U-1995			
	10952997	Oct 10, 2037	U-1995			
	10993941	Oct 10, 2037	U-1995			
	11026931	Aug 14, 2039	U-1995			
	11026939	Sep 18, 2038	DP U-1995			
	11026939	Sep 18, 2038	DP U-3055			
	11040029	Oct 10, 2037	U-1995			
	11311532	Sep 18, 2038	DP U-1995			
	11311532	Sep 18, 2038	DP U-3055			
	11439629	Oct 10, 2037	U-3055			
	11654142	Nov 14, 2038	U-3055			
	8039627	Apr 11, 2031	DS DP			
	8357697	Nov 08, 2027	U-1995			
	8357697	Nov 08, 2027	U-3055			
<u>VALBENAZINE TOSYLATE - INGREZZA</u>						
N 209241 003	10065952	Oct 28, 2036	DS DP U-1995		I-925	Aug 18, 2026
	10065952	Oct 28, 2036	DS DP U-3055		ODE-440	Aug 18, 2030
	10844058	Oct 28, 2036	DS DP U-1995			
	10844058	Oct 28, 2036	DS DP U-3055			
	10851103	Oct 28, 2036	DS DP U-1995			
	10851103	Oct 28, 2036	DS DP U-3055			
	10851104	Oct 28, 2036	DS U-1995			
	10851104	Oct 28, 2036	DS U-3055			
	10857137	Oct 10, 2037	U-1995			
	10857148	Oct 10, 2037	U-1995			
	10874648	Oct 10, 2037	U-1995			
	10874648	Oct 10, 2037	U-3046			
	10874648	Oct 10, 2037	U-3055			
	10906902	Dec 22, 2036	DS DP			
	10906903	Dec 22, 2036	DS DP			
	10912771	Oct 10, 2037	U-1995			
	10912771	Oct 10, 2037	U-3055			
	10912771	Oct 10, 2037	U-3076			
	10919892	Dec 22, 2036	DS DP			
	10940141	Aug 10, 2040	U-1995			
	10952997	Oct 10, 2037	U-1995			
	10993941	Oct 10, 2037	U-1995			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>VALBENAZINE TOSYLATE - INGREZZA</u>						
N 209241 003	11026931	Aug 14, 2039	U-1995			
	11026939	Sep 18, 2038	DP U-1995			
	11026939	Sep 18, 2038	DP U-3055			
	11040029	Oct 10, 2037	U-1995			
	11311532	Sep 18, 2038	DP U-1995			
	11311532	Sep 18, 2038	DP U-3055			
	11439629	Oct 10, 2037	U-3055			
	11654142	Nov 14, 2038	U-3055			
	8039627	Apr 11, 2031	DS DP			
	8357697	Nov 08, 2027	U-1995			
	8357697	Nov 08, 2027	U-3055			
<u>VALBENAZINE TOSYLATE - INGREZZA SPRINKLE</u>						
N 218390 001	10065952	Oct 28, 2036	DS DP U-1995		I-925	Aug 18, 2026
	10065952	Oct 28, 2036	DS DP U-3055			
	10844058	Oct 28, 2036	DS DP U-1995			
	10844058	Oct 28, 2036	DS DP U-3055			
	10851103	Oct 28, 2036	DS DP U-1995			
	10851103	Oct 28, 2036	DS DP U-3055			
	10851104	Oct 28, 2036	DS U-1995			
	10851104	Oct 28, 2036	DS U-3055			
	10857137	Oct 10, 2037	U-1995			
	10857148	Oct 10, 2037	U-1995			
	10874648	Oct 10, 2037	U-1995			
	10874648	Oct 10, 2037	U-3046			
	10874648	Oct 10, 2037	U-3055			
	10906902	Dec 22, 2036	DS DP			
	10906903	Dec 22, 2036	DS DP			
	10912771	Oct 10, 2037	U-1995			
	10912771	Oct 10, 2037	U-3055			
	10912771	Oct 10, 2037	U-3076			
	10919892	Dec 22, 2036	DS DP			
	10940141	Aug 10, 2040	U-1995			
	10952997	Oct 10, 2037	U-1995			
	10993941	Oct 10, 2037	U-1995			
	11026931	Aug 14, 2039	U-1995			
	11026939	Sep 18, 2038	DP U-1995			
	11026939	Sep 18, 2038	DP U-3055			
	11040029	Oct 10, 2037	U-1995			
	11311532	Sep 18, 2038	DP U-1995			
	11311532	Sep 18, 2038	DP U-3055			
	11439629	Oct 10, 2037	U-3055			
	11654142	Nov 14, 2038	U-3055			
	8039627	Apr 11, 2031	DS DP			
	8357697	Nov 08, 2027	U-1995			
	8357697	Nov 08, 2027	U-3055			
<u>VALBENAZINE TOSYLATE - INGREZZA SPRINKLE</u>						
N 218390 002	10065952	Oct 28, 2036	DS DP U-1995		I-925	Aug 18, 2026
	10065952	Oct 28, 2036	DS DP U-3055			
	10844058	Oct 28, 2036	DS DP U-1995			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>VALBENZAZINE TOSYLATE - INGREZZA SPRINKLE</u>						
N 218390 002	10844058	Oct 28, 2036	DS DP U-3055			
	10851103	Oct 28, 2036	DS DP U-1995			
	10851103	Oct 28, 2036	DS DP U-3055			
	10851104	Oct 28, 2036	DS U-1995			
	10851104	Oct 28, 2036	DS U-3055			
	10857137	Oct 10, 2037	U-1995			
	10857148	Oct 10, 2037	U-1995			
	10874648	Oct 10, 2037	U-1995			
	10874648	Oct 10, 2037	U-3046			
	10874648	Oct 10, 2037	U-3055			
	10906902	Dec 22, 2036	DS DP			
	10906903	Dec 22, 2036	DS DP			
	10912771	Oct 10, 2037	U-1995			
	10912771	Oct 10, 2037	U-3055			
	10912771	Oct 10, 2037	U-3076			
	10919892	Dec 22, 2036	DS DP			
	10940141	Aug 10, 2040	U-1995			
	10952997	Oct 10, 2037	U-1995			
	10993941	Oct 10, 2037	U-1995			
	11026931	Aug 14, 2039	U-1995			
	11026939	Sep 18, 2038	DP U-1995			
	11026939	Sep 18, 2038	DP U-3055			
	11040029	Oct 10, 2037	U-1995			
	11311532	Sep 18, 2038	DP U-1995			
	11311532	Sep 18, 2038	DP U-3055			
	11439629	Oct 10, 2037	U-3055			
	11654142	Nov 14, 2038	U-3055			
	8039627	Apr 11, 2031	DS DP			
	8357697	Nov 08, 2027	U-1995			
	8357697	Nov 08, 2027	U-3055			
<u>VALBENZAZINE TOSYLATE - INGREZZA SPRINKLE</u>						
N 218390 003	10065952	Oct 28, 2036	DS DP U-1995		I-925	Aug 18, 2026
	10065952	Oct 28, 2036	DS DP U-3055			
	10844058	Oct 28, 2036	DS DP U-1995			
	10844058	Oct 28, 2036	DS DP U-3055			
	10851103	Oct 28, 2036	DS DP U-1995			
	10851103	Oct 28, 2036	DS DP U-3055			
	10851104	Oct 28, 2036	DS U-1995			
	10851104	Oct 28, 2036	DS U-3055			
	10857137	Oct 10, 2037	U-1995			
	10857148	Oct 10, 2037	U-1995			
	10874648	Oct 10, 2037	U-1995			
	10874648	Oct 10, 2037	U-3046			
	10874648	Oct 10, 2037	U-3055			
	10906902	Dec 22, 2036	DS DP			
	10906903	Dec 22, 2036	DS DP			
	10912771	Oct 10, 2037	U-1995			
	10912771	Oct 10, 2037	U-3055			
	10912771	Oct 10, 2037	U-3076			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>VALBENZAZINE TOSYLATE - INGREZZA SPRINKLE</u>						
N 218390 003	10919892	Dec 22, 2036	DS DP			
	10940141	Aug 10, 2040		U-1995		
	10952997	Oct 10, 2037		U-1995		
	10993941	Oct 10, 2037		U-1995		
	11026931	Aug 14, 2039		U-1995		
	11026939	Sep 18, 2038	DP	U-1995		
	11026939	Sep 18, 2038	DP	U-3055		
	11040029	Oct 10, 2037		U-1995		
	11311532	Sep 18, 2038	DP	U-1995		
	11311532	Sep 18, 2038	DP	U-3055		
	11439629	Oct 10, 2037		U-3055		
	11654142	Nov 14, 2038		U-3055		
	8039627	Apr 11, 2031	DS DP			
	8357697	Nov 08, 2027		U-1995		
	8357697	Nov 08, 2027		U-3055		
<u>VALGANCICLOVIR HYDROCHLORIDE - VALCYTE</u>						
N 022257 001	8889109	Dec 11, 2027	DP			
	9642911	Dec 11, 2027	DP			
<u>VAMOROLONE - AGAMREE</u>						
N 215239 001	10857161	May 28, 2029	DP	U-3747	NCE	Oct 26, 2028
	11382922	Jul 16, 2040	DS DP		ODE-450	Oct 26, 2030
	11471471	Mar 17, 2040		U-3747		
	11690853	Mar 07, 2033		U-3747		
	11833159	May 28, 2029	DP			
	8334279	May 28, 2029		U-3747		
<u>VANCOMYCIN - VANCOMYCIN</u>						
N 213895 001	10039804	Nov 06, 2035	DP	U-282		
	10188697	Nov 06, 2035	DP	U-282		
	10849956	Nov 06, 2035	DP			
	11000567	Nov 06, 2035	DP			
	11517609	Nov 06, 2035		U-282		
<u>VANCOMYCIN HYDROCHLORIDE - FIRVANO KIT</u>						
N 208910 001	10493028	Mar 13, 2035	DP			
	10688046	Mar 13, 2035	DP			
	10959946	Mar 13, 2035	DP			
	10959947	Mar 13, 2035	DP	U-3104		
	10959947	Mar 13, 2035	DP	U-3105		
	10959948	Mar 13, 2035	DP	U-3104		
	10959948	Mar 13, 2035	DP	U-3105		
	10959949	Mar 13, 2035	DP	U-3104		
	10959949	Mar 13, 2035	DP	U-3105		
	11638692	Mar 13, 2035	DP			
<u>VANCOMYCIN HYDROCHLORIDE - FIRVANO KIT</u>						
N 208910 002	10493028	Mar 13, 2035	DP			
	10688046	Mar 13, 2035	DP			
	10959946	Mar 13, 2035	DP			
	10959947	Mar 13, 2035	DP	U-3104		
	10959947	Mar 13, 2035	DP	U-3105		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>VANCOMYCIN HYDROCHLORIDE - FIRVANQ KIT</u>						
N 208910	002	10959948	Mar 13, 2035	DP	U-3104	
		10959948	Mar 13, 2035	DP	U-3105	
		10959949	Mar 13, 2035	DP	U-3104	
		10959949	Mar 13, 2035	DP	U-3105	
		11638692	Mar 13, 2035	DP		
<u>VANCOMYCIN HYDROCHLORIDE - VANCOMYCIN HYDROCHLORIDE</u>						
N 211962	001	10039804	Nov 06, 2035	DP	U-282	
		10188697	Nov 06, 2035	DP	U-282	
		10849956	Nov 06, 2035	DP		
		11517609	Nov 06, 2035		U-282	
		11628200	Nov 06, 2035	DP		
<u>VANCOMYCIN HYDROCHLORIDE - VANCOMYCIN HYDROCHLORIDE</u>						
N 211962	002	10039804	Nov 06, 2035	DP	U-282	
		10188697	Nov 06, 2035	DP	U-282	
		10849956	Nov 06, 2035	DP		
		11517609	Nov 06, 2035		U-282	
		11628200	Nov 06, 2035	DP		
<u>VANCOMYCIN HYDROCHLORIDE - VANCOMYCIN HYDROCHLORIDE</u>						
N 211962	003	10039804	Nov 06, 2035	DP	U-282	
		10188697	Nov 06, 2035	DP	U-282	
		10849956	Nov 06, 2035	DP		
		11517609	Nov 06, 2035		U-282	
		11628200	Nov 06, 2035	DP		
<u>VANCOMYCIN HYDROCHLORIDE - VANCOMYCIN HYDROCHLORIDE</u>						
N 211962	004	10039804	Nov 06, 2035	DP	U-282	
		10188697	Nov 06, 2035	DP	U-282	
		10849956	Nov 06, 2035	DP		
		11517609	Nov 06, 2035		U-282	
		11628200	Nov 06, 2035	DP		
<u>VANCOMYCIN HYDROCHLORIDE - VANCOMYCIN HYDROCHLORIDE</u>						
N 211962	005	10039804	Nov 06, 2035	DP	U-282	
		10188697	Nov 06, 2035	DP	U-282	
		10849956	Nov 06, 2035	DP		
		11517609	Nov 06, 2035		U-282	
		11628200	Nov 06, 2035	DP		
<u>VANCOMYCIN HYDROCHLORIDE - VANCOMYCIN HYDROCHLORIDE</u>						
N 211962	006	10039804	Nov 06, 2035	DP	U-282	
		10188697	Nov 06, 2035	DP	U-282	
		10849956	Nov 06, 2035	DP		
		11517609	Nov 06, 2035		U-282	
		11628200	Nov 06, 2035	DP		
<u>VANCOMYCIN HYDROCHLORIDE - VANCOMYCIN HYDROCHLORIDE</u>						
N 211962	007	10039804	Nov 06, 2035	DP	U-282	
		10188697	Nov 06, 2035	DP	U-282	
		10849956	Nov 06, 2035	DP		
		11517609	Nov 06, 2035		U-282	
		11628200	Nov 06, 2035	DP		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>VANCOMYCIN HYDROCHLORIDE - VANCOMYCIN HYDROCHLORIDE</u>						
N 211962	007	10039804	Nov 06, 2035	DP	U-282	
		10188697	Nov 06, 2035	DP	U-282	
		10849956	Nov 06, 2035	DP		
		11517609	Nov 06, 2035		U-282	
		11628200	Nov 06, 2035	DP		
<u>VANDETANIB - CAPRELSA</u>						
N 022405	001	8067427	Aug 08, 2028	DP		
<u>VANDETANIB - CAPRELSA</u>						
N 022405	002	8067427	Aug 08, 2028	DP		
<u>VARDENAFIL HYDROCHLORIDE - LEVITRA</u>						
N 021400	001	8273876	Jul 23, 2027		U-1288	
<u>VARDENAFIL HYDROCHLORIDE - LEVITRA</u>						
N 021400	002	8273876	Jul 23, 2027		U-1288	
<u>VARDENAFIL HYDROCHLORIDE - LEVITRA</u>						
N 021400	003	8273876	Jul 23, 2027		U-1288	
<u>VARDENAFIL HYDROCHLORIDE - LEVITRA</u>						
N 021400	004	8273876	Jul 23, 2027		U-1288	
<u>VARDENAFIL HYDROCHLORIDE - STAXYN</u>						
N 200179	001	8613950	Dec 23, 2028	DP		
<u>VARENICLINE TARTRATE - TYRVAYA</u>						
N 213978	001	10456396	Oct 19, 2035	DP	U-1900	
		11224598	Oct 19, 2035		U-1900	
		11903941	Oct 19, 2035		U-1900	
		11903942	Oct 19, 2035	DP		
		11903943	Oct 19, 2035		U-1900	
		11911380	Oct 19, 2035		U-1900	
		9504644	Oct 19, 2035		U-1900	
		9504645	Oct 19, 2035	DP		
		9532944	Oct 19, 2035		U-1900	
		9597284	Oct 19, 2035		U-1900	
<u>VASOPRESSIN - VASOSTRICT</u>						
N 204485	001	9375478	Jan 30, 2035		U-1857	
		9687526	Jan 30, 2035		U-1857	
		9744209	Jan 30, 2035		U-1857	
		9744239	Jan 30, 2035		U-1857	
		9750785	Jan 30, 2035	DP		
<u>VASOPRESSIN - VASOSTRICT</u>						
N 204485	002	9375478	Jan 30, 2035		U-1857	
		9687526	Jan 30, 2035		U-1857	
		9744209	Jan 30, 2035		U-1857	
		9744239	Jan 30, 2035		U-1857	
		9750785	Jan 30, 2035	DP		
		9937223	Jan 30, 2035		U-1857	
<u>VASOPRESSIN - VASOSTRICT</u>						
N 204485	003	10010575	Jan 30, 2035		U-1857	

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>VASOPRESSIN - VASOSTRICT</u>						
N 204485	003	9919026	Jan 30, 2035	DP		
		9925233	Jan 30, 2035		U-1857	
		9925234	Jan 30, 2035		U-1857	
		9962422	Jan 30, 2035		U-1857	
		9968649	Jan 30, 2035		U-1857	
		9974827	Jan 30, 2035		U-1857	
		9981006	Jan 30, 2035		U-1857	
<u>VASOPRESSIN - VASOSTRICT</u>						
N 204485	004	10010575	Jan 30, 2035		U-1857	
		9919026	Jan 30, 2035	DP		
		9925233	Jan 30, 2035		U-1857	
		9925234	Jan 30, 2035		U-1857	
		9962422	Jan 30, 2035		U-1857	
		9968649	Jan 30, 2035		U-1857	
		9974827	Jan 30, 2035		U-1857	
		9981006	Jan 30, 2035		U-1857	
<u>VASOPRESSIN - VASOSTRICT</u>						
N 204485	005	10010575	Jan 30, 2035		U-1857	
		9919026	Jan 30, 2035	DP		
		9925233	Jan 30, 2035		U-1857	
		9925234	Jan 30, 2035		U-1857	
		9962422	Jan 30, 2035		U-1857	
		9968649	Jan 30, 2035		U-1857	
		9974827	Jan 30, 2035		U-1857	
		9981006	Jan 30, 2035		U-1857	
<u>VASOPRESSIN - VASOSTRICT</u>						
N 204485	006	10010575	Jan 30, 2035		U-1857	
		9919026	Jan 30, 2035	DP		
		9925233	Jan 30, 2035		U-1857	
		9925234	Jan 30, 2035		U-1857	
		9962422	Jan 30, 2035		U-1857	
		9968649	Jan 30, 2035		U-1857	
		9974827	Jan 30, 2035		U-1857	
		9981006	Jan 30, 2035		U-1857	
<u>VEMURAFENIB - ZELBORAF</u>						
N 202429	001	7504509	Oct 22, 2026	DS DP		
		7863288	Jun 20, 2029	DS DP		
		8143271	Jun 21, 2026	DS DP		
		8470818	Aug 02, 2026		U-1418	
		8470818	Aug 02, 2026		U-2164	
		8741920	Jul 27, 2030	DS DP		
		9447089	Jun 06, 2032	DP		
<u>VENETOCLAX - VENCLEXTA</u>						
N 208573	001	10730873	Nov 21, 2031	DS	ODE-185	Jun 08, 2025
		10993942	Sep 06, 2033		ODE-211	Nov 21, 2025
		11110087	Sep 06, 2033		ODE-239	May 15, 2026
		11110087	Sep 06, 2033		U-3223	
		11369599	May 23, 2032	DP		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>VENETOCLAX - VENCLEXTA</u>						
N 208573	001	11413282	Sep 06, 2033	U-3412		
		11590128	Sep 06, 2033	U-3548		
		8546399	Jun 27, 2031	DS DP		
		8722657	Jan 29, 2032	DS		
		9174982	May 26, 2030	U-2323		
		9174982	May 26, 2030	U-2445		
		9174982	May 26, 2030	U-2446		
		9174982	May 26, 2030	U-2537		
		9539251	Sep 06, 2033	U-2538		
<u>VENETOCLAX - VENCLEXTA</u>						
N 208573	002	10730873	Nov 21, 2031	DS	ODE-185	Jun 08, 2025
		10993942	Sep 06, 2033	U-3114	ODE-211	Nov 21, 2025
		11110087	Sep 06, 2033	U-3222	ODE-239	May 15, 2026
		11110087	Sep 06, 2033	U-3223		
		11369599	May 23, 2032	DP		
		11413282	Sep 06, 2033	U-3412		
		11590128	Sep 06, 2033	U-3548		
		8546399	Jun 27, 2031	DS DP		
		8722657	Jan 29, 2032	DS		
		9174982	May 26, 2030	U-2323		
		9174982	May 26, 2030	U-2445		
		9174982	May 26, 2030	U-2446		
		9174982	May 26, 2030	U-2537		
		9539251	Sep 06, 2033	U-2538		
<u>VENETOCLAX - VENCLEXTA</u>						
N 208573	003	10730873	Nov 21, 2031	DS	ODE-185	Jun 08, 2025
		10993942	Sep 06, 2033	U-3114	ODE-211	Nov 21, 2025
		11110087	Sep 06, 2033	U-3222	ODE-239	May 15, 2026
		11110087	Sep 06, 2033	U-3223		
		11369599	May 23, 2032	DP		
		11413282	Sep 06, 2033	U-3412		
		11590128	Sep 06, 2033	U-3548		
		8546399	Jun 27, 2031	DS DP		
		8722657	Jan 29, 2032	DS		
		9174982	May 26, 2030	U-2323		
		9174982	May 26, 2030	U-2445		
		9174982	May 26, 2030	U-2446		
		9174982	May 26, 2030	U-2537		
		9539251	Sep 06, 2033	U-2538		
<u>VENLAFAXINE BESYLATE - VENLAFAXINE BESYLATE</u>						
N 215429	001	7776358	May 16, 2028	DP		
<u>VERICIGUAT - VEROUVO</u>						
N 214377	001	10736896	May 19, 2031	DS DP	NCE	Jan 19, 2026
		11439642	May 19, 2031	U-3062		
		8420656	May 19, 2031	DS DP		
		8921377	May 19, 2031	U-3062		
		9604948	Nov 26, 2032	DS DP	U-3062	
		9993476	May 19, 2031	U-3062		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>VERICIGUAT - VEROUVO</u>						
N 214377	001	10736896	May 19, 2031	DS DP	NCE	Jan 19, 2026
		11439642	May 19, 2031	U-3062		
		8420656	May 19, 2031	DS DP		
		8921377	May 19, 2031	U-3062		
		9604948	Nov 26, 2032	DS DP U-3062		
		9993476	May 19, 2031	U-3062		
<u>VERICIGUAT - VEROUVO</u>						
N 214377	002	10736896	May 19, 2031	DS DP	NCE	Jan 19, 2026
		11439642	May 19, 2031	U-3062		
		8420656	May 19, 2031	DS DP		
		8921377	May 19, 2031	U-3062		
		9604948	Nov 26, 2032	DS DP U-3062		
		9993476	May 19, 2031	U-3062		
<u>VERICIGUAT - VEROUVO</u>						
N 214377	003	10736896	May 19, 2031	DS DP	NCE	Jan 19, 2026
		11439642	May 19, 2031	U-3062		
		8420656	May 19, 2031	DS DP		
		8921377	May 19, 2031	U-3062		
		9604948	Nov 26, 2032	DS DP U-3062		
		9993476	May 19, 2031	U-3062		
<u>VIBEGRON - GEMTESA</u>						
N 213006	001	12102638	Mar 22, 2040	U-3045	NCE	Dec 23, 2025
		8247415	Dec 01, 2030	DS DP U-3045		
		8653260	Apr 02, 2029	DS		
<u>VIGABATRIN - VIGAFYDE</u>						
N 217684	001	12016857	Aug 16, 2039	DP		
<u>VILOXAZINE HYDROCHLORIDE - QELBREE</u>						
N 211964	001	11324753	Sep 04, 2029	U-727	NCE	Apr 02, 2026
		11458143	Sep 04, 2029	U-727	NPP	Apr 29, 2025
		12121523	Sep 04, 2029	U-727		
		9358204	Feb 07, 2033	DP		
		9603853	Feb 07, 2033	U-727		
		9662338	Feb 07, 2033	DP		
<u>VILOXAZINE HYDROCHLORIDE - QELBREE</u>						
N 211964	002	11324753	Sep 04, 2029	U-727	NCE	Apr 02, 2026
		11458143	Sep 04, 2029	U-727	NPP	Apr 29, 2025
		12121523	Sep 04, 2029	U-727		
		9358204	Feb 07, 2033	DP		
		9603853	Feb 07, 2033	U-727		
		9662338	Feb 07, 2033	DP		
<u>VILOXAZINE HYDROCHLORIDE - QELBREE</u>						
N 211964	003	11324753	Sep 04, 2029	U-727	NCE	Apr 02, 2026
		11458143	Sep 04, 2029	U-727	NPP	Apr 29, 2025
		12121523	Sep 04, 2029	U-727		
		9358204	Feb 07, 2033	DP		
		9603853	Feb 07, 2033	U-727		
		9662338	Feb 07, 2033	DP		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>VILOXAZINE HYDROCHLORIDE - QELBREE</u>						
N 211964	003 11324753	Sep 04, 2029	U-727		NCE	Apr 02, 2026
	11458143	Sep 04, 2029	U-727		NPP	Apr 29, 2025
	12121523	Sep 04, 2029	U-727			
	9358204	Feb 07, 2033	DP			
	9603853	Feb 07, 2033	U-727			
	9662338	Feb 07, 2033	DP			
<u>VILTOLARSEN - VILTEPSO</u>						
N 212154	001 10870676	Aug 31, 2031	DS DP U-3039		NCE	Aug 12, 2025
	9079934	Aug 31, 2031	DS DP		ODE-280	Aug 12, 2027
<u>VISMODEGIB - ERIVEDGE</u>						
N 203388	001 7888364	Nov 11, 2028	DS DP			
	9278961	Dec 15, 2028	U-1825			
	9790183	Sep 02, 2025	U-3109			
<u>VOCLOSPORIN - LUPKYNIS</u>						
N 213716	001 10286036	Dec 07, 2037	U-3056		NCE	Jan 22, 2026
	11622991	Dec 07, 2037	U-3056			
	7332472	Oct 17, 2025	DS DP U-3056			
<u>VONOPRAZAN FUMARATE - VOQUEZNA</u>						
N 215151	001 7977488	Aug 11, 2028	DS		I-948	Jul 17, 2027
	9186411	Aug 11, 2030	DP		NCE	May 03, 2027
					NP	Nov 01, 2026
<u>VONOPRAZAN FUMARATE - VOQUEZNA</u>						
N 215151	002 7977488	Aug 11, 2028	DS		NCE	May 03, 2027
	9186411	Aug 11, 2030	DP		NP	Nov 01, 2026
<u>VORAPAXAR SULFATE - ZONTIVITY</u>						
N 204886	001 7304078	Dec 23, 2027	DS DP U-1512			
<u>VORASIDENIB - VORANIGO</u>						
N 218784	001 10172864	Jul 11, 2034	DS DP		NCE	Aug 06, 2029
	11345677	Jan 16, 2039	DS DP U-3978		ODE-491	Aug 06, 2031
	11844758	Dec 04, 2035	U-3977			
	9579324	Jul 11, 2034	DS			
<u>VORASIDENIB - VORANIGO</u>						
N 218784	002 10172864	Jul 11, 2034	DS DP		NCE	Aug 06, 2029
	11345677	Jan 16, 2039	DS DP U-3978		ODE-491	Aug 06, 2031
	11844758	Dec 04, 2035	U-3977			
	9579324	Jul 11, 2034	DS			
<u>VORINOSTAT - ZOLINZA</u>						
N 021991	001 7399787	Feb 09, 2025	U-892			
	7456219	Mar 11, 2027	DS			
	8093295	May 16, 2026	DP			
	8450372	Mar 18, 2028	U-892			
<u>VORTIOXETINE HYDROBROMIDE - TRINTELLIX</u>						
N 204447	001 11458134	Jun 15, 2027	DP U-3463		M-232	Aug 23, 2026
	11458134*PED	Dec 15, 2027			PED	Feb 23, 2027
	7144884	Jun 17, 2026	DS DP U-1439			
	7144884*PED	Dec 17, 2026				

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>VORTIOXETINE HYDROBROMIDE - TRINTELLIX</u>						
N 204447 001	8722684	Jun 30, 2031	DS DP			
	8722684*PED	Dec 30, 2031				
	8969355	Jun 15, 2027		U-1668		
	8969355*PED	Dec 15, 2027				
	9125908	Jun 15, 2027		U-2309		
	9125908*PED	Dec 15, 2027				
	9125909	Jun 15, 2027		U-2309		
	9125909*PED	Dec 15, 2027				
	9125910	Jun 15, 2027		U-2309		
	9125910*PED	Dec 15, 2027				
	9227946	Jun 15, 2027		U-1668		
	9227946*PED	Dec 15, 2027				
	9278096	Mar 21, 2032		U-2436		
	9278096*PED	Sep 21, 2032				
	9861630	Jun 15, 2027		U-1668		
	9861630*PED	Dec 15, 2027				
<u>VORTIOXETINE HYDROBROMIDE - TRINTELLIX</u>						
N 204447 002	11458134	Jun 15, 2027	DP U-3463		M-232	Aug 23, 2026
	11458134*PED	Dec 15, 2027			PED	Feb 23, 2027
	7144884	Jun 17, 2026	DS DP U-1439			
	7144884*PED	Dec 17, 2026				
	8722684	Jun 30, 2031	DS DP			
	8722684*PED	Dec 30, 2031				
	8969355	Jun 15, 2027		U-1668		
	8969355*PED	Dec 15, 2027				
	9125908	Jun 15, 2027		U-2309		
	9125908*PED	Dec 15, 2027				
	9125909	Jun 15, 2027		U-2309		
	9125909*PED	Dec 15, 2027				
	9125910	Jun 15, 2027		U-2309		
	9125910*PED	Dec 15, 2027				
	9227946	Jun 15, 2027		U-1668		
	9227946*PED	Dec 15, 2027				
	9278096	Mar 21, 2032		U-2436		
	9278096*PED	Sep 21, 2032				
	9861630	Jun 15, 2027		U-1668		
	9861630*PED	Dec 15, 2027				
<u>VORTIOXETINE HYDROBROMIDE - TRINTELLIX</u>						
N 204447 003	11458134	Jun 15, 2027	DP U-3463		M-232	Aug 23, 2026
	11458134*PED	Dec 15, 2027			PED	Feb 23, 2027
	7144884	Jun 17, 2026	DS DP U-1439			
	7144884*PED	Dec 17, 2026				
	8722684	Jun 30, 2031	DS DP			
	8722684*PED	Dec 30, 2031				
	8969355	Jun 15, 2027		U-1668		
	8969355*PED	Dec 15, 2027				
	9125908	Jun 15, 2027		U-2309		
	9125908*PED	Dec 15, 2027				
	9125909	Jun 15, 2027		U-2309		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>VORTIOXETINE HYDROBROMIDE - TRINTELLIX</u>						
N 204447 003	9125909*PED	Dec 15, 2027				
	9125910	Jun 15, 2027	U-2309			
	9125910*PED	Dec 15, 2027				
	9227946	Jun 15, 2027	U-1668			
	9227946*PED	Dec 15, 2027				
	9278096	Mar 21, 2032	U-2436			
	9278096*PED	Sep 21, 2032				
	9861630	Jun 15, 2027	U-1668			
	9861630*PED	Dec 15, 2027				
<u>VORTIOXETINE HYDROBROMIDE - TRINTELLIX</u>						
N 204447 004	11458134	Jun 15, 2027	DP U-3463		M-232	Aug 23, 2026
	11458134*PED	Dec 15, 2027			PED	Feb 23, 2027
	7144884	Jun 17, 2026	DS DP U-1439			
	7144884*PED	Dec 17, 2026				
	8722684	Jun 30, 2031	DS DP			
	8722684*PED	Dec 30, 2031				
	8969355	Jun 15, 2027	U-1668			
	8969355*PED	Dec 15, 2027				
	9125908	Jun 15, 2027	U-2309			
	9125908*PED	Dec 15, 2027				
	9125909	Jun 15, 2027	U-2309			
	9125909*PED	Dec 15, 2027				
	9125910	Jun 15, 2027	U-2309			
	9125910*PED	Dec 15, 2027				
	9227946	Jun 15, 2027	U-1668			
	9227946*PED	Dec 15, 2027				
	9278096	Mar 21, 2032	U-2436			
	9278096*PED	Sep 21, 2032				
	9861630	Jun 15, 2027	U-1668			
	9861630*PED	Dec 15, 2027				
<u>VOSORITIDE - VOXZOGO</u>						
N 214938 001	10646550	Aug 01, 2036	DP U-3927		NCE	Nov 19, 2026
	11590204	Aug 01, 2036	DP U-3927		NPP	Oct 20, 2026
	11911446	Aug 01, 2036	U-3927		ODE-387	Nov 19, 2028
	8198242	Jun 11, 2030	DS DP U-3927		ODE-449	Oct 20, 2030
	9907834	Aug 01, 2036	DP			
	RE48267	May 20, 2030	U-3927			
<u>VOSORITIDE - VOXZOGO</u>						
N 214938 002	10646550	Aug 01, 2036	DP U-3927		NCE	Nov 19, 2026
	11590204	Aug 01, 2036	DP U-3927		NPP	Oct 20, 2026
	11911446	Aug 01, 2036	U-3927		ODE-387	Nov 19, 2028
	8198242	Jun 11, 2030	DS DP U-3927		ODE-449	Oct 20, 2030
	9907834	Aug 01, 2036	DP			
	RE48267	May 20, 2030	U-3927			
<u>VOSORITIDE - VOXZOGO</u>						
N 214938 003	10646550	Aug 01, 2036	DP U-3927		NCE	Nov 19, 2026
	11590204	Aug 01, 2036	DP U-3927		NPP	Oct 20, 2026
	11911446	Aug 01, 2036	U-3927		ODE-387	Nov 19, 2028

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>VOXORITIDE - VOXZOGO</u>						
N 214938	003 8198242	Jun 11, 2030	DS DP U-3927		ODE-449	Oct 20, 2030
	9907834	Aug 01, 2036	DP			
	RE48267	May 20, 2030	U-3927			
<u>VOXELOTOR - OXBRYTA</u>						
N 213137	001 10017491	Dec 28, 2032	DP		ODE-281	Nov 25, 2026
	10034879	Dec 28, 2032	DS DP		ODE-394	Dec 17, 2028
	10493035	Oct 12, 2037	DP			
	10722502	Feb 06, 2035	DP			
	10806733	Dec 28, 2032	DS			
	11020382	Dec 02, 2036	U-3133			
	11020382	Dec 02, 2036	U-3134			
	11452720	Feb 06, 2035	U-3459			
	11944612	Dec 02, 2036	U-3893			
	9018210	Nov 25, 2033	DS DP			
	9248199	Jan 29, 2034	U-2676			
	9248199	Jan 29, 2034	U-2715			
	9447071	Feb 06, 2035	DS DP			
<u>VOXELOTOR - OXBRYTA</u>						
N 213137	002 10017491	Dec 28, 2032	DP		ODE-394	Dec 17, 2028
	10034879	Dec 28, 2032	DS DP			
	10493035	Oct 12, 2037	DP			
	10722502	Feb 06, 2035	DP			
	10806733	Dec 28, 2032	DS			
	11020382	Dec 02, 2036	U-3133			
	11020382	Dec 02, 2036	U-3134			
	11452720	Feb 06, 2035	U-3459			
	11944612	Dec 02, 2036	U-3893			
	9018210	Nov 25, 2033	DS DP			
	9248199	Jan 29, 2034	U-2676			
	9248199	Jan 29, 2034	U-2715			
	9447071	Feb 06, 2035	DS DP			
<u>VOXELOTOR - OXBRYTA</u>						
N 216157	001 10017491	Dec 28, 2032	DP		ODE-394	Dec 17, 2028
	10034879	Dec 28, 2032	DS DP			
	10722502	Feb 06, 2035	DP			
	10806733	Dec 28, 2032	DS			
	11020382	Dec 02, 2036	U-3133			
	11020382	Dec 02, 2036	U-3134			
	11452720	Feb 06, 2035	U-3459			
	11944612	Dec 02, 2036	U-3893			
	9018210	Nov 25, 2033	DS DP			
	9248199	Jan 29, 2034	U-2676			
	9248199	Jan 29, 2034	U-2715			
	9447071	Feb 06, 2035	DS DP			
<u>VUTRISIRAN SODIUM - AMVUTTRA</u>						
N 215515	001 10131907	Aug 24, 2028	DS DP U-3396		NCE	Jun 13, 2027
	10208307	Jul 28, 2036	DS DP U-3396		ODE-212	Jun 13, 2029
	10570391	Nov 16, 2032	DS DP U-3396			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>VUTRISIRAN SODIUM - AMVUTTRA</u>						
N 215515 001	10612024	Aug 14, 2035	DS DP U-3396			
	10683501	Jul 28, 2036	DS DP U-3396			
	10806791	Dec 04, 2028	DS			
	11286486	Jul 28, 2036	DS DP U-3396			
	11401517	Aug 14, 2035	DS DP U-3396			
	12049628	Jul 28, 2036	DS DP U-3396			
	8106022	Dec 12, 2029	DS DP U-3396			
	8828956	Dec 04, 2028	DS DP U-3396			
	9370581	Dec 04, 2028	DS DP U-3396			
	9399775	Nov 16, 2032	DS DP U-3396			
<u>XENON XE-129 HYPERPOLARIZED - XENOVUE</u>						
N 214375 001	10583205	Feb 20, 2035	DP		NCE	Dec 23, 2027
	11052161	Dec 29, 2035	DP			
<u>ZANUBRUTINIB - BRUKINSA</u>						
N 213217 001	10570139	Apr 22, 2034	U-1745		I-817	Jan 19, 2026
	10570139	Apr 22, 2034	U-2145		I-936	Mar 07, 2027
	10570139	Apr 22, 2034	U-2537		ODE-274	Jan 19, 2030
	10570139	Apr 22, 2034	U-2666		ODE-276	Nov 14, 2026
	10570139	Apr 22, 2034	U-3063		ODE-370	Sep 14, 2028
	10570139	Apr 22, 2034	U-3486		ODE-371	Aug 31, 2028
	10570139	Apr 22, 2034	U-3860		ODE-467	Mar 07, 2031
	10927117	Aug 15, 2037	DS DP			
	11142528	Apr 22, 2034	DP U-1745			
	11142528	Apr 22, 2034	DP U-2145			
	11142528	Apr 22, 2034	DP U-2537			
	11142528	Apr 22, 2034	DP U-2666			
	11142528	Apr 22, 2034	DP U-3063			
	11142528	Apr 22, 2034	DP U-3486			
	11142528	Apr 22, 2034	DP U-3860			
	11591340	Aug 15, 2037	U-1745			
	11591340	Aug 15, 2037	U-2145			
	11591340	Aug 15, 2037	U-2537			
	11591340	Aug 15, 2037	U-2666			
	11591340	Aug 15, 2037	U-3063			
	11591340	Aug 15, 2037	U-3486			
	11591340	Aug 15, 2037	U-3727			
	11591340	Aug 15, 2037	U-3728			
	11591340	Aug 15, 2037	U-3729			
	11701357	Jun 24, 2039	U-3860			
	11786531	Jan 19, 2043	U-3715			
	11786531	Jan 19, 2043	U-3716			
	11786531	Jan 19, 2043	U-3717			
	11786531	Jan 19, 2043	U-3718			
	11786531	Jan 19, 2043	U-3719			
	11786531	Jan 19, 2043	U-3720			
	11786531	Jan 19, 2043	U-3875			
	11851437	Aug 15, 2037	DS DP			
	11884674	Aug 15, 2037	U-1745			
	11884674	Aug 15, 2037	U-2145			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>ZANUBRUTINIB - BRUKINSA</u>						
N 213217 001	11884674	Aug 15, 2037		U-2537		
	11884674	Aug 15, 2037		U-2666		
	11884674	Aug 15, 2037		U-3063		
	11884674	Aug 15, 2037		U-3486		
	11884674	Aug 15, 2037		U-3860		
	11896596	Jan 19, 2043		U-3715		
	11896596	Jan 19, 2043		U-3716		
	11896596	Jan 19, 2043		U-3717		
	11896596	Jan 19, 2043		U-3718		
	11896596	Jan 19, 2043		U-3719		
	11896596	Jan 19, 2043		U-3720		
	11896596	Jan 19, 2043		U-3875		
	11911386	Jan 19, 2043		U-3715		
	11911386	Jan 19, 2043		U-3716		
	11911386	Jan 19, 2043		U-3717		
	11911386	Jan 19, 2043		U-3718		
	11911386	Jan 19, 2043		U-3719		
	11911386	Jan 19, 2043		U-3720		
	11911386	Jan 19, 2043		U-3875		
	11970500	Aug 15, 2037		U-1745		
	11970500	Aug 15, 2037		U-2145		
	11970500	Aug 15, 2037		U-2537		
	11970500	Aug 15, 2037		U-2666		
	11970500	Aug 15, 2037		U-3063		
	11970500	Aug 15, 2037		U-3486		
	11970500	Aug 15, 2037		U-3860		
	9447106	Apr 22, 2034	DS DP	U-1745		
	9447106	Apr 22, 2034	DS DP	U-2145		
	9447106	Apr 22, 2034	DS DP	U-2537		
	9447106	Apr 22, 2034	DS DP	U-2666		
	9447106	Apr 22, 2034	DS DP	U-3063		
	9447106	Apr 22, 2034	DS DP	U-3486		
	9447106	Apr 22, 2034	DS DP	U-3860		
<u>ZAVEGEPANT HYDROCHLORIDE - ZAVZPRET</u>						
N 216386 001	8481546	Oct 07, 2031	DS DP	U-3555	NCE	Mar 09, 2028
<u>ZILUCOPLAN SODIUM - ZILBRYSO</u>						
N 216834 001	10106579	Jun 12, 2035	DS DP	U-3735	NCE	Oct 17, 2028
	10208089	Jun 12, 2035	DS	U-3735	ODE-446	Oct 17, 2030
	10435438	Jun 12, 2035	DS	U-3735		
	10562934	Jun 12, 2035	DS			
	10835574	Jun 12, 2035		DP		
	11014965	Jun 12, 2035	DS	U-3735		
	11535650	Jun 12, 2035		DP		
	11752190	Jun 12, 2035		DP U-3735		
	11965040	Jun 12, 2035		DP		
<u>ZILUCOPLAN SODIUM - ZILBRYSO</u>						
N 216834 002	10106579	Jun 12, 2035	DS DP	U-3735	NCE	Oct 17, 2028
	10208089	Jun 12, 2035	DS	U-3735	ODE-446	Oct 17, 2030

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>ZILUCOPLAN SODIUM - ZILBRYSO</u>						
N 216834 002	10435438	Jun 12, 2035	DS U-3735			
	10562934	Jun 12, 2035	DS			
	10835574	Jun 12, 2035	DP			
	11014965	Jun 12, 2035	DS U-3735			
	11535650	Jun 12, 2035	DP			
	11752190	Jun 12, 2035	DP U-3735			
	11965040	Jun 12, 2035	DP			
<u>ZILUCOPLAN SODIUM - ZILBRYSO</u>						
N 216834 003	10106579	Jun 12, 2035	DS DP U-3735		NCE	Oct 17, 2028
	10208089	Jun 12, 2035	DS U-3735		ODE-446	Oct 17, 2030
	10435438	Jun 12, 2035	DS U-3735			
	10562934	Jun 12, 2035	DS			
	10835574	Jun 12, 2035	DP			
	11014965	Jun 12, 2035	DS U-3735			
	11535650	Jun 12, 2035	DP			
	11752190	Jun 12, 2035	DP U-3735			
	11965040	Jun 12, 2035	DP			
<u>ZINC CHLORIDE - ZINC CHLORIDE</u>						
A 216152 001					CGT	Apr 14, 2025
<u>ZIPRASIDONE HYDROCHLORIDE - GEODON</u>						
N 020825 001					M-232	Jan 28, 2025
<u>ZIPRASIDONE HYDROCHLORIDE - GEODON</u>						
N 020825 002					M-232	Jan 28, 2025
<u>ZIPRASIDONE HYDROCHLORIDE - GEODON</u>						
N 020825 003					M-232	Jan 28, 2025
<u>ZIPRASIDONE HYDROCHLORIDE - GEODON</u>						
N 020825 004					M-232	Jan 28, 2025
<u>ZOLEDRONIC ACID - ZOMETA</u>						
N 021223 002	8324189	May 29, 2025	U-1308			
	8324189	May 29, 2025	U-1309			
	8324189	May 29, 2025	U-53			
<u>ZOLEDRONIC ACID - ZOMETA</u>						
N 021223 003	7932241	Feb 05, 2028	DP			
	8324189	May 29, 2025	U-1308			
	8324189	May 29, 2025	U-1309			
	8324189	May 29, 2025	U-53			
<u>ZOLEDRONIC ACID - RECLAST</u>						
N 021817 001	7932241	Feb 05, 2028	DP			
<u>ZOLPIDEM TARTRATE - EDLUAR</u>						
N 021997 001	9265720	Feb 25, 2031	U-674			
	9597281	Apr 06, 2027	U-674			
<u>ZOLPIDEM TARTRATE - EDLUAR</u>						
N 021997 002	9265720	Feb 25, 2031	U-674			
	9597281	Apr 06, 2027	U-674			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>ZOLPIDEM TARTRATE - ZOLPIMIST</u>						
N 022196	001 8236285	Aug 07, 2032	DS DP U-70			
<u>ZOLPIDEM TARTRATE - INTERMEZZO</u>						
N 022328	001 7658945	Apr 15, 2027	DP U-1194			
	7682628	Feb 16, 2025	U-1194			
	8242131	Aug 20, 2029	U-1266			
	8252809	Feb 16, 2025	DP			
<u>ZOLPIDEM TARTRATE - INTERMEZZO</u>						
N 022328	002 7658945	Apr 15, 2027	DP U-1194			
	7682628	Feb 16, 2025	U-1194			
	8242131	Aug 20, 2029	U-1266			
	8252809	Feb 16, 2025	DP			
<u>ZONISAMIDE - ZONISADE</u>						
N 214273	001 11478456	Aug 18, 2038	U-3458			
	11529333	Aug 18, 2038	DP			
<u>ZURANOLONE - ZURZUVAE</u>						
N 217369	001 10172871	Apr 17, 2034	U-2552		NCE	Oct 31, 2028
	10342810	Apr 17, 2034	U-2552			
	11236121	Aug 23, 2037	DS			
	11884696	Dec 24, 2037	U-2552			
	9512165	Apr 17, 2034	DS DP			
<u>ZURANOLONE - ZURZUVAE</u>						
N 217369	002 10172871	Apr 17, 2034	U-2552		NCE	Oct 31, 2028
	10342810	Apr 17, 2034	U-2552			
	11236121	Aug 23, 2037	DS			
	11884696	Dec 24, 2037	U-2552			
	9512165	Apr 17, 2034	DS DP			
<u>ZURANOLONE - ZURZUVAE</u>						
N 217369	003 10172871	Apr 17, 2034	U-2552		NCE	Oct 31, 2028
	10342810	Apr 17, 2034	U-2552			
	11236121	Aug 23, 2037	DS			
	11884696	Dec 24, 2037	U-2552			
	9512165	Apr 17, 2034	DS DP			

PATENT AND EXCLUSIVITY TERMS

ADB 1 of 225

PATENT & EXCLUSIVITY ABBREVIATIONS

CGT	COMPETITIVE GENERIC THERAPY
D	NEW DOSING SCHEDULE (SEE INDIVIDUAL REFERENCES)
GAIN	GAIN EXCLUSIVITY
I	NEW INDICATION (SEE INDIVIDUAL REFERENCES)
M	MISCELLANEOUS EXCLUSIVITY CODES (SEE INDIVIDUAL REFERENCES)
NC	NEW COMBINATION
NCE	NEW CHEMICAL ENTITY
NCE*	NEW CHEMICAL ENTITY (AN ENANTIOMER OF PREVIOUSLY APPROVED RACEMIC MIXTURE. SEE SECTION 505(U) OF THE FEDERAL FOOD AND DRUG COSMETIC ACT).
NDF	NEW DOSAGE FORM
NE	NEW ESTER OR SALT OF AN ACTIVE INGREDIENT
NP	NEW PRODUCT
NP*	NEW PRODUCT (MINT FLAVORED)
NPP	NEW PATIENT POPULATION
NR	NEW ROUTE
NS	NEW STRENGTH
ODE	ORPHAN DRUG EXCLUSIVITY (SEE INDIVIDUAL REFERENCES)
ODE*	FDA has not recognized Orphan-Drug Exclusivity (ODE) for this drug, but it contains the same active moiety or moieties as another drug(s) that was eligible for ODE, and also shares ODE-protected use(s) or indication(s) with that drug(s). An application seeking approval for the same active moiety or moieties, including an ANDA that cites this NDA as its basis of submission, may not be approved for such ODE-protected use(s) and indication(s)
PC	PATENT CHALLENGE
PED	PEDIATRIC EXCLUSIVITY
RTO	RX TO OTC SWITCH OR OTC USE
RTO*	OTC USE FOR WOMEN AGES 15 AND 16
RTO**	OTC USE FOR WOMEN 14 AND BELOW
U	PATENT USE CODE (SEE INDIVIDUAL REFERENCES)
W	EXCLUSIVITY ON THIS APPLICATION EXPIRING ON THIS DATE HAS BEEN WAIVED BY SPONSOR - SEE SECTION 1.8 OF ORANGE BOOK PREFACE WAIVED EXCLUSIVITY

EXCLUSIVITY DOSING SCHEDULE

D-1	ONCE A DAY APPLICATION
D-2	ONCE DAILY DOSING
D-3	SEVEN DAYS/SEVEN DAYS/SEVEN DAYS DOSING SCHEDULE
D-4	SEVEN DAYS/FOURTEEN DAYS DOSING SCHEDULE
D-5	TEN DAYS/ELEVEN DAYS DOSING SCHEDULE
D-6	SEVEN DAYS/NINE DAYS/FIVE DAYS DOSING SCHEDULE
D-7	BID DOSING
D-8	INTRAVENOUS, EPIDURAL AND INTRATHECAL DOSING
D-9	NARCOTIC OVERDOSE IN ADULTS
D-10	NARCOTIC OVERDOSE IN CHILDREN
D-11	POSTOPERATIVE NARCOTIC DEPRESSION IN CHILDREN
D-12	BEDTIME DOSING OF 800MG FOR TREATMENT OF ACTIVE DUODENAL ULCER
D-13	INCREASED MAXIMUM DAILY DOSAGE RECOMMENDATION
D-14	BEDTIME DOSING OF 800MG FOR TREATMENT OF ACTIVE BENIGN GASTRIC ULCER
D-15	SINGLE DAILY DOSE OF 25MG/37.5MG
D-16	CONTINUOUS INTRAVENOUS INFUSION
D-17	400MG EVERY 12 HOURS FOR THREE DAYS FOR UNCOMPLICATED URINARY TRACT INFECTIONS
D-18	LOWER RECOMMENDED STARTING DOSE GUIDELINES
D-19	BOLUS DOSING GUIDELINES
D-20	SINGLE 32MG DOSE
D-21	ALTERNATIVE DOSAGE OF 300MG ONCE DAILY AFTER THE EVENING MEAL

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY DOSING SCHEDULE**

- D-22 REDUCTION IN INFUSION TIME FROM 24 TO 4 HOURS FOR THE 60MG DOSE
- D-23 INCREASE MAXIMUM DOSE AND VARIATIONS IN THE DOSING REGIMEN
- D-24 FOR OVARIAN CANCER THE RECOMMENDED REGIMEN IS 135MG/M2 OR 175MG/M2 INTRAVENOUSLY OVER THREE HOURS EVERY THREE WEEKS
- D-25 ADDITIONAL DOSAGE REGIMEN EQUAL TO HALF THE ORIGINAL DOSING REGIMEN
- D-26 ONCE WEEKLY APPLICATION
- D-27 BID DOSING IN PATIENTS 12 YEARS OF AGE AND OLDER FOR PREVENTION OF NAUSEA AND VOMITING ASSOCIATED WITH MODERATE EMETOGENIC CANCER CHEMOTHERAPY
- D-28 USE OF ISOVUE-370 IN EXCRETORY UROGRAPHY AT EQUIVALENT GRAMS OF IODINE TO THE CURRENTLY APPROVED ISOVUE-250 AND ISOVUE-300
- D-29 INCREASE OF CUMULATIVE DOSE TO 0.3MMOL/KG FOR MRI OF CNS IN ADULTS
- D-30 5000 IU DOSE FOR PROPHYLAXIS AGAINST DEEP VEIN THROMBOSIS
- D-31 CHANGE IN RECOMMENDED TOTAL DAILY DOSE TO 80MG (40MG BID)
- D-32 REMOVAL OF THE RESTRICTIONS LIMITING TREATMENT TO TWO CONSECUTIVE WEEKS AND TO SMALL AREAS
- D-33 ONCE DAILY DOSING FOR PLAQUE PSORIASIS
- D-34 EVERY FOUR MONTHS DOSAGE REGIMEN
- D-35 FOR A ONE WEEK DOSING OF INTERDIGITAL TINEA PEDIS
- D-36 FOR A SINGLE 2MG DOSE AS AN ALTERNATIVE TO THE 1MG DOSE GIVEN TWICE DAILY
- D-37 DOSING REGIMEN FOR ADMINISTRATION EITHER ONCE DAILY (QD) OR TWICE DAILY (BID)
- D-38 CONTINUOUS INFUSION AS AN ALTERNATE METHOD OF ADMINISTRATION
- D-39 CHANGE IN TIME TO TAKE THE DRUG PRIOR TO A MEAL TO PREVENT MEAL-INDUCED HEARTBURN SYMPTOMS FROM ".1/2 TO 1 HOUR BEFORE EATING" TO ".. RIGHT BEFORE EATING OR UP TO 60MIN BEFORE CONSUMING..."
- D-40 ONCE-A-DAY DOSING REGIMEN
- D-41 DRUG MAY BE DOSED RIGHT BEFORE A MEAL OR ANY TIME UP TO 30MIN BEFORE EATING OR DRINKING FOOD AND BEVERAGES THAT WOULD BE EXPECTED TO CAUSE SYMPTOMS
- D-42 TEN DAY DOSING REGIMEN FOR TRIPLE THERAPY, PREVACID IN COMBINATION WITH CLARITHROMYCIN AND AMOXICILLIN, FOR THE ERADICATION OF H.PYLORI IN PATIENTS WITH DUODENAL ULCER DISEASE
- D-43 INITIATION OF TREATMENT WITH 900MG/DAY BY DELETION OF THE REQUIREMENT TO TITRATE TO 900MG/DAY OVER A 3-DAY PERIOD
- D-44 IN A CLINICAL TRIAL, FEWER DISCONTINUATIONS DUE TO ADVERSE EVENTS, ESPECIALLY DIZZINESS AND VERTIGO, WERE OBSERVED WHEN TITRATING THE DOSE IN INCREMENTS OF 50MG/DAY EVERY 3 DAYS UNTIL AN EFFECTIVE DOSE (NOT EXCEEDING 400MG/DAY) WAS REACHED
- D-45 ONCE DAILY DOSING FOR MAINTENANCE ONLY
- D-46 NEW DOSING REGIMEN OF 80MG DAILY
- D-47 PREVENTION OF HEARTBURN SYMPTOMS WHEN ADMINISTERED FROM 15 MINUTES UP TO, BUT NOT INCLUDING, 1 HOUR PRIOR TO A PROVOCATIVE MEAL
- D-48 ADMINISTRATION OF CISATRACURIUM A NEUROMUSCULAR BLOCKING AGENT AT DOSES OF 3 AND 4X THE ED95 OF CISATRACURIUM FOLLOWING INDUCTION WITH THIOPENTAL
- D-49 PEDIATRIC DOSING GUIDELINES
- D-50 INFORMATION FOR USE OF CORVERT IN POST-CARDIAC SURGERY PATIENTS
- D-51 OPTIONAL STARTING DOSE OF 40MG/DAY
- D-52 ALTERNATE DOSING REGIMEN OF 1250MG TWICE DAILY
- D-53 USE IN PEDIATRIC PATIENTS FROM 1 MONTH TO 16 YEARS OF AGE
- D-54 USE OF ZYBAN FOR MAINTENANCE THERAPY. TREATMENT UP TO 6 MONTHS WAS SHOWN EFFICACIOUS
- D-55 ADDITION OF A HIGHER DOSE OF NUTROPIN FOR PUBERTAL PATIENTS (PUBERTAL DOSE LESS THAN OR EQUAL TO 0.7MG/KG/WEEK)
- D-56 ADDITION OF POSTPRANDIAL DOSING

PATENT AND EXCLUSIVITY TERMS

ADB 3 of 225

EXCLUSIVITY DOSING SCHEDULE

- D-57 3-HOUR INFUSION OF TAXOL GIVEN EVERY THREE WEEKS AT A DOSE OF 175MG/M2 FOLLOWED BY CISPLATIN AT A DOSE OF 75MG/M2 FOR THE FIRST-LINE TREATMENT OF ADVANCED OVARIAN CANCER
- D-58 CHANGE IN DOSING INTERVAL TO ONCE-DAILY ADMINISTRATION
- D-59 REDUCTION OF ELEVATED LDL-C IN A NEW, HIGHER STRENGTH TABLET, 0.8MG, AND FOR EXTENSION OF THE DOSAGE RANGE TO 0.8MG DAILY
- D-60 ADDITION OF A POST-OPERATIVE DOSING REGIMEN
- D-61 ONCE WEEKLY DOSING FOR THE TREATMENT OF POSTMENOPAUSAL OSTEOPOROSIS
- D-62 ONCE WEEKLY DOSING FOR THE PREVENTION OF POSTMENOPAUSAL OSTEOPOROSIS
- D-63 TO ALLOW A TITRATION DOSING REGIMEN USING A 25MG DOSE
- D-64 INCREASING DOSAGE FOR NERVE BLOCK ANESTHESIA USING NAROPIN 7.5MG/ML AND FOR EXTENDING THE DURATION OF TREATMENT FOR POSTOPERATIVE ANALGESIA USING NAROPIN 2MG/ML
- D-65 CHANGE DOSING AND ADMINISTRATION TO INDICATE MAINTENANCE OF WEIGHT LOSS OVER AN 18 MONTH PERIOD THUS EXTENDING THE USE OF THIS DRUG FROM ONE TO TWO YEARS
- D-66 DOSING RECOMMENDATIONS FOR PATIENTS UNDERGOING PCI
- D-67 SHORTER TREATMENT COURSE OF THREE DAYS IN THE TREATMENT OF RECURRENT EPISODES OF GENITAL HERPES
- D-68 CHANGE OF ADMIN RATE FOR INFUSION OF AREDIA FOR TREATMENT OF MODERATE AND SEVERE HYPERCALCEMIA OF MALIGNANCY FROM 24 HOURS TO 2 HOURS UP TO BUT NOT INCLUDING 24 HOURS
- D-69 SHORTENED DOSING REGIMEN TO 5 DAYS FOR THE TREATMENT OF ACUTE EXACERBATION OF CHRONIC BRONCHITIS
- D-70 80MG ONCE DAILY DOSING REGIMEN
- D-71 EIGHT WEEK DOSING REGIMEN
- D-72 INFORMATION REGARDING INCREASED RATE OF INFUSION FOR DEPACON
- D-73 ONCE A WEEK DOSING FOR THE TREATMENT OF POSTMENOPAUSAL OSTEOPOROSIS
- D-74 ONCE A WEEK DOSING FOR THE PREVENTION OF POSTMENOPAUSAL OSTEOPOROSIS
- D-75 INTERMITTENT DOSING REGIMEN, STARTING DAILY DOSE 14 DAYS PRIOR TO THE ANTICIPATED ONSET OF MENSTRUATION THROUGH THE FIRST FULL DAY OF MENSES AND REPEATING WITH EACH NEW CYCLE
- D-76 FOR USE ON AN "AS NEEDED" OR PRN BASIS FOR THE MANAGEMENT OF NASAL SYMPTOMS IN PATIENTS FOR WHOM THE DRUG IS INDICATED
- D-77 ADDITION OF 20MG AND 40MG DAILY AS OPTIONAL STARTING DOSES WITH 40MG INTENDED FOR PATIENTS WHO REQUIRE A LARGE REDUCTION IN LDL-C (MORE THAN 45%)
- D-78 USE OF FLEXERIL 5MG FOR THE RELIEF OF MUSCLE SPASM ASSOCIATED WITH ACUTE, PAINFUL, MUSCULOSKELETAL CONDITIONS
- D-79 NEW LOWER STARTING DOSE FOR TREATMENT OF MODERATE TO SEVERE VASOMOTOR SYMPTOMS AND/OR MODERATE TO SEVERE SYMPTOMS OF VULVAR AND VAGINAL ATROPHY ASSOCIATED W/ THE MENOPAUSE
- D-80 CHANGE OF DOSING SCHEDULE FOR LANTUS FROM ONCE DAILY AT BEDTIME TO FLEXIBLE DAILY DOSING
- D-81 NEW LOWER STARTING DOSE FOR THE PREVENTION OF POSTMENOPAUSAL OSTEOPORSIS
- D-82 USE OF PREMARIN 0.3 MG AND 0.45 MG FOR THE PREVENTION OF POSTMENOPAUSAL OSTEOPOROSIS
- D-83 750 MG, ONCE DAILY FOR 5 DAYS FOR COMMUNITY ACQUIRED PNEUMONIA (CAP)
- D-84 ONCE-A-DAY DOSING OF FLOXACIN OTIC FOR THE TREATMENT OF ADULTS AND PEDIATRIC PATIENTS (AGES 6 MO & OLDER) W/ OTITIS EXTERNA CAUSED BY SUSCEPTIBLE STRAINS OF E.COLI, P.AERUGINOSA AND S.AUREUS
- D-85 LOWER RECOMMENDED STARTING DOSE GUIDELINES FOR TREATMENT OF MODERATE TO SEVERE VASOMOTOR SYMPTOMS ASSOCIATED WITH THE MENOPAUSE
- D-86 FOR USE IN SELECT EXTERNAL INSULIN PUMPS
- D-87 ADDITION OF ONCE-WEEKLY DOSING FOR THE TREATMENT TO INCREASE BONE MASS IN MEN WITH OSTEOPOROSIS

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY DOSING SCHEDULE**

- D-88 NEW DOSING RANGE OF 200-400MG PER DAY IN TWO DIVIDED DOSES FOR ADULTS WITH PARTIAL SEIZURES
- D-89 USE OF REYATAZ 300 MG/RITONAVIR 100 MG ONCE DAILY FOR TREATMENT IN HIV-INFECTED ANTIRETROVIRAL-EXPERIENCED PATIENTS
- D-90 ADDITION OF DAYTIME ADMINISTRATION TO TREAT VULVOVAGINAL CANDIDIASIS
- D-91 ALTERNATE INTERMITTENT DOSING REGIMEN
- D-92 ALTERNATIVE DOSAGE OF 1000MG ONCE DAILY AT BEDTIME
- D-93 ALTERNATE TWO OR THREE TIMES DAILY DOSING REGIMENS
- D-94 NEW MAXIMUM DOSAGE OF 72 MG/DAY IN ADOLESCENTS 13-17 YEARS OF AGE WITH ATTENTION DEFICIT HYPERACTIVITY DISORDER (ADHD)
- D-95 BROADENED INITIAL STARTING DOSE FOR HYPERTENSION FROM 50 MG TO 100 MG TO 25 MG TO 100 MG DOSE RANGE
- D-96 ONCE-MONTHLY TREATMENT OF POSTMENOPAUSAL OSTEOPOROSIS WITH BONIVA (IBANDRONATE SODIUM) 150 MG TABLETS
- D-97 PED CANCER PT POPULATION EXPANDED TO INCLUDE PTS 6 MOS UP TO BUT NOT INCLUDING 4 YRS AND DOSING INSTRUCTIONS TO ADMIN 30 MIN BEFORE CHEMO WITH SECOND AND THIRD DOSES 4 & 8 HOURS AFTER FIRST DOSE
- D-98 DOSING FOR PED SURGICAL PTS EXPANDED TO INCLUDE PTS 1 MONTH UP TO BUT NOT INCLUDING 2 YEARS OF AGE
- D-99 ONCE DAILY ADMINISTRATION FOR THE TREATMENT OF HIV INFECTION IN THERAPY NAIVE ADULT PATIENTS
- D-100 750 MG ONCE DAILY FOR FIVE DAYS FOR THE TREATMENT OF ACUTE BACTERIAL SINUSITIS
- D-101 ONCE DAILY IN CHRONIC IDIOPATHIC URTICARIA FOR ADULTS AND CHILDREN 12 YEARS OF AGE AND OLDER
- D-102 NEW DOSING REGIMEN OF ONE SPRAY TWICE DAILY FOR SEASONAL ALLERGIC RHINITIS IN PATIENTS 12 YRS OF AGE AND OLDER
- D-103 NEW DOSING RECOMMENDATION FOR THE TREATMENT OF RECURRENT GENITAL HERPES IN IMMUNOCOMPETENT PATIENTS, SPECIFICALLY A REDUCTION IN COURSE OF THERAPY FROM FAMCICLOVIR 125 MG TWICE-A-DAY FOR 5 DAYS TO 1000 MG TWICE-A-DAY FOR 1 DAY.
- D-104 0.5MG/0.1MG FOR THE TREATMENT OF MODERATE TO SEVERE VASOMOTOR SYMPTOMS ASSOCIATED WITH MENOPAUSE IN WOMEN WHO HAVE A UTERUS
- D-105 USE OF ACTONEL 75MG TWO CONSECUTIVE DAYS PER MONTH FOR THE PREVENTION AND TREATMENT OF POSTMENOPAUSAL OSTEOPOROSIS
- D-106 FIVE DAY TREATMENT OF SELECTED SUSCEPTIBLE STRAINS OF STREPTOCOCCUS PNEUMONIAE, HAEMOPHILUS INFLUENZA, MYCOPLASMA PNEUMONIAE, AND CHLAMYDIA PNEUMONIAE FOR COMMUNITY-ACQUIRED PNEUMONIA
- D-107 PROVIDES FOR THE COMBINATION TABLET OF 70MG ALENDRONATE AND 5600 IU OF VITAMIN D3 FOR THE TREATMENT OF OSTEOPOROSIS IN POSTMENOPAUSAL WOMEN AND TO INCREASE BONE MASS IN MEN WITH OSTEOPOROSIS
- D-108 TREATMENT OF COMPLICATED URINARY TRACT INFECTION AND ACUTE PYELONEPHRITIS WITH LEVAQUIN 750MG ONCE DAILY FOR FIVE DAYS
- D-109 PROVIDE FOR THE USE OF A LOWER DOSE FOR THE TREATMENT OF ADULTS WITH CHRONIC PHASE CHRONIC MYELOID LEUKEMIA (CML) WITH RESISTANCE OR INTOLERANCE TO PRIOR THERAPY INCLUDING IMATINIB MESYLATE
- D-110 TREATMENT OF SCHIZOPHRENIA IN ADOLESCENTS AGED 13-17
- D-111 PROVIDES FOR ONCE DAILY USE OF CIALIS, 2.5 MG AND 5 MG, FOR THE TREATMENT OF ERECTILE DYSFUNCTION
- D-112 PROVIDES FOR PEDIATRIC PUMP USE
- D-113 ONCE DAILY DOSING REGIMEN FOR PATIENTS WHO BECOME CONSTIPATED ON TWICE DAILY REGIMEN
- D-114 NEW DOSING RECOMMENDATIONS FOR USE OF SIROLIMUS IN COMBINATION WITH CYCLOSPORINE FOR THE PROPHYLAXIS OF REJECTION IN HIGH-RISK RENAL TRANSPLANT RECIPIENTS
- D-115 STARTING DOSE OF 15MG/DAY FOR MONOTHERAPY IN ACUTE TREATMENT OF BIPOLAR DISORDER, MANIC OR MIXED
- D-116 ALTERNATIVE DOSING REGIMEN ATAZANAVIR SULFATE CO-ADMINISTERED WITH RITONAVIR FOR THE TREATMENT OF HIV-1 INFECTION IN TREATMENT NAIVE PATIENTS

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY DOSING SCHEDULE**

D-117	50 MG TABLET FOR INITIATION OF DOSE TITRATION FOR BIPOLAR DISORDER
D-118	TWO 400MG TABLETS ONCE DAILY, CO-ADMINISTERED WITH 100MG RITONAVIR
D-119	DOSING RECOMMENDATIONS FOR HIV INFECTED PEDIATRIC PATIENTS 6 TO LESS THAN 18 YEARS OF AGE
D-120	DOSING REGIMEN ADJUSTMENTS
D-121	CHANGE TO REMOVE 20 MG MAXIMUM DOSAGE RESTRICTION
D-122	USE OF VAGIFEM 10 MCG FOR THE TREATMENT OF ATROPHIC VAGINITIS DUE TO MENOPAUSE
D-123	ALTERNATIVE DOSING REGIMEN DOSE OF 20 MG/METER SQUARE BY CONTINUOUS INTRAVENOUS INFUSION OVER 1 HOUR REPEATED DAILY FOR 5 DAYS
D-124	ONCE DAILY DOSING REGIMEN IN ADULT PATIENTS WITH LESS THAN THREE LOPINAVIR RESISTANCE-ASSOCIATED SUBSTITUTIONS
D-125	EXTEND CURRENT DOSING REGIMEN TO 900MG (2-450MG TABLETS) ONCE A DAY WITHIN 10 DAYS OF TRANSPLANTATION UNTIL 200 DAYS POST-TRANSPLANTATION FOR THE PREVENTION OF CYTOMEGALOVIRUS (CMV) DISEASE IN ADULT KIDNEY TRANSPLANT PATIENTS AT HIGH RISK.
D-126	CHANGE DOSAGE REGIMEN FROM 250MG TO 500MG
D-127	DOSING REGIMEN FOR ADULT PATIENTS WITH CHRONIC HEPATITIS B (CHB) AND DECOMPENSATED LIVER DISEASE
D-128	SINGLE IV DOSE OF FOSAPREPITANT 150MG, DOSED CONCOMITANTLY WITH 5HT3 RECEPTOR ANTAGONIST & CORTICOSTEROID, FOR PREVENTION OF ACUTE & DELAYED NAUSEA & VOMITING ASSOCIATED WITH INITIAL AND REPEAT COURSES OF HIGHLY EMETOGENIC CANCER CHEMO
D-129	800/100 MG DARUNAVIR/RITONAVIR, ONCE DAILY, IN TREATMENT-EXPERIENCED HIV-1 INFECTED PATIENTS WITH NO DARUNAVIR RESISTANCE ASSOCIATED SUBSTITUTIONS
D-130	DOSING RECOMMENDATIONS FOR TREATMENT OF HIV-1 INFECTION DURING PREGNANCY BASED ON DATA FROM STUDY AI424-182, A STUDY OF ATAZANAVIR/RITONAVIR IN COMBINATION WITH ZIDOVUDINE/LAMIVUDINE IN HIV INFECTED PREGNANT WOMEN
D-131	EVERY 6 TO 8 WEEKS FOR THE 120MG STRENGTH FOR PATIENTS WHO ARE CONTROLLED ON SOMATULINE DEPOT 60MG OR 90MG
D-132	45MG FOR 6 MONTH ADMINISTRATION
D-133	NEW EFFICACY DATA AND DOSING REGIMEN FOR PREGNANCY IN NORMAL OVULATORY WOMEN UNDERGOING CONTROLLED OVARIAN STIMULATION AS PART OF AN IVF OR INTRACYTOPLASMIC SPERM INJECTION (ICSI) CYCLE
D-134	INCREASING MAXIMUM DOSING OF PATIENTS WITH SCHIZOPHRENIA TO 160 MG/DAY
D-135	UPDATE LABELING WITH ONCE DAILY DOSING IN HIV-1 INFECTED, TREATMENT-NAIVE PEDIATRIC PATIENTS 12 TO LESS THAN 18 YEARS OF AGE
D-136	ALTERNATE DOSING REGIMEN FOR UNCOMPLICATED URETHRAL OR ENDOCERVICAL INFECTION CAUSED BY CHLAMYDIA TRACHOMATIS, ADMINISTER 200 MG BY MOUTH ONCE-A-DAY FOR 7 DAYS
D-137	NEW LOWER DOSING REGIMEN FOR REVATIO IN THE TREATMENT OF PULMONARY ARTERIAL HYPERTENSION (WHO GROUP 1) IN ADULTS
D-138	80 MG DOSING REGIMEN FOR THE RISK REDUCTION OF REBLEEDING OF GASTRIC AND DUODENAL ULCERS IN THE FIRST 72 HOURS FOLLOWING THERAPEUTIC ENDOSCOPY IN ADULTS
D-139	ADDITIONAL INFORMATION ADDED TO THE DOSING AND ADMINISTRATION SECTION OF THE LABELING REGARDING THE ADMINISTRATION OF BRAVELLE AND MENOPUR IN THE SAME SYRINGE TO OVULATORY WOMEN AS PART OF AN ART CYCLE
D-140	REVISED DOSING SCHEDULE TO ADMINISTER AVANAFIL 15 MINUTES PRIOR TO SEXUAL ACTIVITY
D-141	DOSING INFORMATION IN PREVIOUSLY UNTREATED MANTLE CELL LYMPHOMA
D-142	DOSE MODIFICATION GUIDELINES FOR BORTEZOMIB WHEN GIVEN IN COMBINATION WITH RITUXIMAB, CYCLOPHOSPHAMIDE, DOXORUBICIN, AND PREDNISONE
D-143	INITIATION OF VIMPAT THERAPY WITH A LOADING DOSE OF 200MG
D-144	LOWER LIMIT OF 15 MINUTES FOR THE INFUSION DURATION
D-145	UPDATES TO THE DOSAGE AND ADMINISTRATION SECTION OF THE LABELING TO REFLECT THE RESULTS OF TWO SHORT TERM STUDIES EVALUATING THE SAFETY AND EFFICACY OF INTUNIV IN CHILDREN AND ADOLESCENTS AGES 6 TO 17 WITH ADHD.

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY DOSING SCHEDULE**

D-146	CHANGE IN TARGET DOSING TO 20MG TO 40MG ORALLY ONCE DAILY
D-147	ONCE DAILY DOSING IN PEDIATRIC PATIENTS 3 MONTHS OF AGE AND OLDER IN COMBINATION WITH OTHER ANTIRETROVIRAL AGENTS FOR THE TREATMENT OF HIV-1 INFECTION
D-148	EXTENDED THE DURATION OF THE DOSING REGIMEN FROM 100 DAYS TO 200 DAYS POST-TRANSPLANTATION FOR THE PREVENTION OF CMV DISEASE IN PEDIATRIC KIDNEY TRANSPLANT
D-149	DOSING INFORMATION ADDED TO THE LABELING REGARDING PEDIATRIC PATIENTS 6 YEARS AND OLDER WITH ITP
D-150	1600MG DAILY FOR PATIENTS ON ADJUNCTIVE THERAPY WHO DID NOT ACHIEVE A SATISFACTORY RESPONSE ON 1200MG DAILY DOSE
D-151	DOSING RECOMMENDATIONS FOR THE TREATMENT OF CHRONIC HEPATITIS C IN PATIENTS CO-INFECTED WITH HIV-1
D-152	DOSING RECOMMENDATIONS AS NECESSARY FOR FEVER AND PAIN FOR AGES 6MO TO LESS THAN 12 YEARS AND 12 TO 17 YEARS.
D-153	IN COMBINATION WITH RIBAVIRIN FOR 12 WEEKS, FOR THE TREATMENT OF GENOTYPE 1, CHRONIC HEPATITIS C TREATMENT EXPERIENCED PATIENTS WITH COMPENSATED CIRRHOSIS BASED UPON THE RESULTS OF THE SIRIUS STUDY
D-154	ADDITION OF A 1500MG-SINGLE-DOSE REGIMEN FOR THE TREATMENT OF ADULT PATIENTS WITH ACUTE BACTERIAL SKIN AND SKIN STRUCTURE INFECTIONS (ABSSI)
D-155	SINGLE IV DOSE OF FOSAPREPITANT 150MG DOSED CONCOMITANTLY WITH 5HT3 RECEPTOR ANTAGONIST & CORTICOSTEROID FOR PREVENTION OF DELAYED NAUSEA AND VOMITING ASSOCIATED WITH INITIAL AND REPEAT COURSES OF MODERATELY EMETOGENIC CANCER CHEMOTHERAPY
D-156	DOSING INFORMATION ADDED TO THE LABELING PROVIDING INFORMATION ON TRANSITIONING FROM SUBCUTANEOUS OR INTRAVENOUS ROUTES OF ADMINISTRATION OF TREPROSTINIL
D-157	UPDATED INFORMATION ADDED TO THE DOSAGE AND ADMINISTRATION SECTION OF THE LABELING PROVIDING DOSAGE RECOMMENDATIONS FOR INTERRUPTIONS AND DISCONTINUATION OF THERAPY
D-158	REVISED DOSING TO EXPAND PATIENT POPULATION TO INCLUDE LIVER TRANSPLANT RECIPIENTS WITH GENOTYPE 1 HCV INFECTION
D-159	REVISED DOSING TO EXPAND PATIENT POPULATION TO INCLUDE LIVER TRANSPLANT RECIPIENTS WITH GENOTYPE 4 HCV INFECTION
D-160	REVISED DOSING TO EXPAND PATIENT POPULATION TO INCLUDE PATIENTS WITH DECOMPENSATED CIRRHOSIS WITH GENOTYPE 1 HCV INFECTION
D-161	DOSAGE RECOMMENDATIONS ADDED TO INCLUDE TREATMENT OF HCV GENOTYPE 3 SUBJECTS CO-INFECTED WITH HIV-1
D-162	DOSING TO INCLUDE PATIENTS WITH CHRONIC HCV GENOTYPE 1 INFECTION WITH COMPENSATED (CHILD-PUGH A) OR DECOMPENSATED (CHILD-PUGH B OR C) CIRRHOSIS AND TREATMENT OF CHRONIC HCV GENOTYPE 3 INFECTION IN SUBJECTS WITH DECOMPENSATED (CHILD-PUGH B OR C) CIRRHOSIS
D-163	DOSING TO INCLUDE PATIENTS WITH CHRONIC HCV GENOTYPE 1A INFECTION WITH COMPENSATED (CHILD-PUGH A) CIRRHOSIS AND GENOTYPE 1B WITH OR WITHOUT COMPENSATED (CHILD-PUGH A) CIRRHOSIS
D-164	UPDATES TO THE DOSAGE AND ADMINISTRATION, DOSE MODIFICATIONS SECTION OF THE LABELING
D-165	DOSING RECOMMENDATION ADDED TO THE LABELING FOR IMBRUVICA USE IN COMBINATION WITH BENDAMUSTINE AND RITUXIMAB FOR THE TREATMENT OF CHRONIC LYMPHOCYTIC LEUKEMIA (CLL)/SMALL LYMPHOCYTIC LEUKEMIA (SLL)
D-166	BROADEN INITIAL STARTING DOSE FOR BIPOLAR I DISORDER TO 5-10MG TWICE DAILY
D-167	ADDITION OF 1200 MG ONCE DAILY DOSING FOR TREATMENT-NAIVE PATIENTS OR PATIENTS WHO ARE VIROLOGICALLY SUPPRESSED ON AN INITIAL REGIMEN OF RALTEGRAVIR FILM-COATED TABLETS 400 MG TWICE DAILY
D-168	NEW DOSING REGIMEN OF 10 MG ONCE DAILY FOR THE REDUCTION IN THE RISK OF RECURRENCE OF DEEP VEIN THROMBOSIS (DVT) AND/OR PULMONARY EMBOLISM (PE) IN PATIENTS AT CONTINUED RISK FOR DVT AND/OR PE AFTER COMPLETION OF INITIAL TREATMENT LASTING AT LEAST 6 MONTHS
D-169	ONCE-DAILY DOSING FOR PATIENTS 5 YEARS OF AGE AND OLDER WHO HAVE UNDETECTABLE SERUM AND URINE SUCCINYLACETONE CONCENTRATIONS AFTER A MINIMUM OF 4 WEEKS ON A STABLE DOSAGE OF NITISINONE

PATENT AND EXCLUSIVITY TERMS

ADB 7 of 225

EXCLUSIVITY DOSING SCHEDULE

- D-170 TO ALLOW WITHDRAWAL THERAPY OF PATIENTS WITH PHILADELPHIA CHROMOSOME POSITIVE CHRONIC MYELOID LEUKEMIA IN CHRONIC PHASE WHO HAVE ACHIEVED A SUSTAINED MOLECULAR RESPONSE ON NILOTINIB THERAPY FOR A MINIMUM OF ONE YEAR PRIOR TO DISCONTINUATION
- D-171 REVISED DOSING TO INCLUDE UP-TITRATION AS A STRATEGY TO IMPROVE TOLERABILITY AND THEREBY REDUCE TREATMENT DISCONTINUATION FOR ROFLUMILAST MAINTENANCE DOSAGE OF 500 MCG DAILY
- D-172 ADDITION OF A ONCE WEEKLY DOSING REGIMEN FOR CARFILZOMIB IN COMBINATION WITH DEXAMETHASONE FOR PATIENTS WITH RELAPSED OR REFRACTORY MULTIPLE MYELOMA WHO HAVE RECEIVED ONE TO THREE LINES OF THERAPY
- D-173 DOSING RECOMMENDATION FOR THE USE OF ELVITEGRAVIR/COBICISTAT/EMTRICITABINE/TENOFOVIR ALAFENAMIDE FIXED DOSE COMBINATION IN HIV-1 INFECTED ADULT PATIENTS WITH END-STAGE-RENAL DISEASE WHO ARE RECEIVING CHRONIC HEMODIALYSIS
- D-174 MODIFICATIONS TO THE EXISTING DOSING REGIMEN TO ALLOW FOR TREATMENT INTERRUPTIONS OF UP TO 8 WEEKS FOR INTOLERABLE ADVERSE REACTIONS
- D-175 EIGHT-WEEK DOSING REGIMEN FOR THE TREATMENT OF GENOTYPES 1, 2, 3, 4, 5, AND 6, CHRONIC HEPATITIS C VIRUS INFECTION IN TREATMENT-NAIVE SUBJECTS WITH COMPENSATED CIRRHOSIS BASED ON THE RESULTS FROM THE EXPEDITION-8 STUDY
- D-176 IBRUTINIB IN COMBINATION WITH RITUXIMAB
- D-177 INFORMATION ADDED TO THE DOSING SECTION IN REGARD TO THE TREATMENT OF CHRONIC HEPATITIS C VIRUS INFECTION IN PATIENTS WITH SEVERE RENAL IMPAIRMENT INCLUDING PATIENTS WITH END STAGE RENAL DISEASE ON DIALYSIS
- D-181 DOSING REGIMEN EXTENDING THE CONTRACEPTION USE FROM 5 YEARS TO UP TO 6 YEARS
- D-182 NEW DOSING REGIMEN FOR THE PREVENTION AND MANAGEMENT OF NERATINIB ASSOCIATED DIARRHEA
- D-183 3-DAY DOSING REGIMEN FOR THE TREATMENT OF COVID-19 IN ADULTS AND PEDIATRIC PATIENTS (>12 YEARS AND WEIGHING AT LEAST 40 KG) WITH POSITIVE RESULTS OF DIRECT SARS-COV-2 VIRAL TESTING, WHO ARE NOT HOSPITALIZED AND HAVE MILD-TO-MODERATE COVID-19, AND ARE AT HIGH RISK FOR PROGRESSION TO SEVERE COVID-19, INCLUDING HOSPITALIZATION OR DEATH
- D-184 NEW DOSING SCHEDULE FOR CABOTEGRAVIR/RILPIVRINE INJECTION EVERY 2 MONTHS
- D-185 ADDITION OF A 3RD MAINTENANCE DOSE OF SEMAGLUTIDE
- D-186 ADDITION OF A 3-DAY FOSAPREPITANT FOR INJECTION INTRAVENOUS DOSING REGIMEN IN PEDIATRIC PATIENTS FOR THE CURRENTLY APPROVED PREVENTION OF CHEMOTHERAPY-INDUCED NAUSEA AND VOMITING
- D-187 ADDITION OF SECOND DOSE FOR TREATMENT OF PRESBYOPIA IN ADULTS
- D-188 USE OF ALTERNATE INITIAL DOSING REGIMEN FOR INITIATION OF LACOSAMIDE TREATMENT IN PARTIAL ONSET SEIZURE PATIENTS ≥ 1 MONTH TO < 17 YEARS OF AGE AND IN PRIMARY GENERALIZED TONIC-CLONIC SEIZURE PATIENTS ≥ 4 TO < 17 YEARS
- D-189 EXTENSION OF LETERMOVIR DOSING REGIMEN FROM 100 TO 200 DAYS POST-TRANSPLANT FOR THE PROPHYLAXIS OF CYTOMEGALOVIRUS (CMV) INFECTION AND DISEASE IN ADULT CMV SEROPOSITIVE RECIPIENTS (R+) OF AN ALLOGENEIC HEMATOPOIETIC STEM CELL TRANSPLANT WHO ARE AT RISK FOR LATE CMV INFECTION AND DISEASE
- D-190 USE OF SEMAGLUTIDE 1.7 MG SUBCUTANEOUS WEEKLY AS AN ADDITIONAL MAINTENANCE DOSE
- D-191 ADDITION OF A ONCE DAILY DOSING REGIMEN
- D-192 EXPANSION OF THE TREATMENT FIELD ON THE FACE OR SCALP UP TO 100 CM²
- D-193 NEW INSERTER INTENDED TO FACILITATE SINGLE-HAND PLACEMENT
- D-194 INCREASE IN THE APPROVED DOSAGE REGIMEN TO A MAXIMUM APPLICATION AREA OF 60 CM² IN TOTAL (CORRESPONDING TO THREE 2 GM TUBES)

EXCLUSIVITY INDICATION

- I-1 DYSMENORRHEA
- I-2 CHOLANGIOPANCREATOGRAPHY
- I-3 INTRAVENOUS DIGITAL SUBTRACTION ANGIOGRAPHY

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

I-4	PERIPHERAL VENOGRAPHY (PHLEBOGRAPHY)
I-5	HYSTEOSALPINGOGRAPHY
I-6	TREATMENT OF JUVENILE ARTHRITIS
I-7	BIOPSY PROVEN MINIMAL CHANGE NEPHROTIC SYNDROME IN CHILDREN
I-8	ADULT INTRAVENOUS CONTRAST-ENHANCED COMPUTED TOMOGRAPHY OF THE HEAD AND BODY
I-9	PREVENTION OF POSTOPERATIVE NAUSEA AND VOMITING
I-10	PREVENTION OF POSTOPERATIVE DEEP VENOUS THROMBOSIS AND PULMONARY EMBOLISM IN TOTAL HIP REPLACEMENT SURGERY
I-11	RELIEF OF MILD TO MODERATE PAIN
I-12	TREATMENT OF CUTANEOUS CANDIDIASIS
I-13	URINARY TRACT INFECTION (UTI) PREVENTION FOR PERIODS UP TO FIVE MONTHS IN WOMEN WITH A HISTORY OF RECURRENT UTI
I-14	SEBORRHEIC DERMATITIS
I-15	PHOTOPHERESIS IN THE PALLIATIVE TREATMENT OF SKIN MANIFESTATIONS OF CUTANEOUS T-CELL LYMPHOMA IN PERSONS NOT RESPONSIVE TO OTHER TREATMENT
I-16	STIMULATE THE DEVELOPMENT OF MULTIPLE FOLLICLES/OOCYTES IN OVULATORY PATIENTS PARTICIPATING IN AN IN VITRO FERTILIZATION PROGRAM
I-17	MANAGEMENT OF CONGESTIVE HEART FAILURE
I-18	ENDOSCOPIC RETROGRADE PANCREATOGRAPHY
I-19	HERNIOGRAPHY
I-20	KNEE ARTHROGRAPHY
I-21	HIGH DOSE METHOTREXATE WITH LEUCOVORIN RESCUE IN COMBINATION WITH OTHER CHEMOTHERAPEUTIC AGENTS TO DELAY RECURRENCE IN PATIENTS WITH NONMETASTATIC OSTEOSARCOMA WHO HAVE UNDERGONE SURGICAL RESECTION OR AMPUTATION FOR THE PRIMARY TUMOR
I-22	RESCUE AFTER HIGH-DOSE METHOTREXATE THERAPY IN OSTEOSARCOMA
I-23	SHORT-TERM TREATMENT OF ACTIVE BENIGN GASTRIC ULCER
I-24	TREATMENT OF RHEUMATOID ARTHRITIS
I-25	ADULT INTRA-ARTERIAL DIGITAL SUBTRACTION ANGIOGRAPHY OF THE HEAD, NECK, ABDOMINAL, RENAL AND PERIPHERAL VESSELS
I-26	TREATMENT OF LIVER FLUKES
I-27	ADJUNCTIVE THERAPY TO DIET TO REDUCE THE RISK OF CORONARY ARTERY DISEASE
I-28	SELECTIVE ADULT VISCERAL ARTERIOGRAPHY
I-29	METASTATIC BREAST CANCER IN PREMENOPAUSAL WOMEN AS AN ALTERNATIVE TO OOPHORECTOMY OR OVARIAN IRRADIATION
I-30	TREATMENT OF TINEA PEDIS
I-31	CONTRAST ENHANCEMENT AGENT TO FACILITATE VISUALIZATION OF LESIONS IN THE SPINE AND ASSOCIATED TISSUES
I-32	PEDIATRIC MYELOGRAPHY
I-33	ORAL USE OF DILUTED OMNIPAQUE INJECTION IN ADULTS FOR CONTRAST ENHANCED COMPUTED TOMOGRAPHY OF THE ABDOMEN
I-34	ORAL USE IN ADULTS FOR PASS-THROUGH EXAMINATION OF THE GASTROINTESTINAL TRACT
I-35	PEDIATRIC CONTRAST ENHANCEMENT OF COMPUTED TOMOGRAPHIC HEAD IMAGING
I-36	ARTHROGRAPHY OF THE SHOULDER JOINTS IN ADULTS
I-37	RADIOGRAPHY OF THE TEMPOROMANDIBULAR JOINT IN ADULTS
I-38	CONTRAST ENHANCEMENT AGENT TO FACILITATE VISUALIZATION OF LESIONS OF THE CENTRAL NERVOUS SYSTEM IN CHILDREN (2 YEARS OF AGE AND OLDER)
I-39	TREATMENT OF ACUTE MYOCARDIAL INFARCTION
I-40	PRIMARY NOCTURNAL ENURESIS
I-41	MIGRAINE HEADACHE PROPHYLAXIS

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

I-42 HERPES ZOSTER

I-43 HERPES SIMPLEX ENCEPHALITIS

I-44 MAINTENANCE THERAPY IN HEALED DUODENAL ULCER PATIENTS AT DOSE OF 1 GRAM TWICE DAILY

I-45 ACUTE TREATMENT OF VARICELLA ZOSTER VIRUS

I-46 USE IN PEDIATRIC COMPUTED TOMOGRAPHIC HEAD AND BODY IMAGING

I-47 TREATMENT OF PEDIATRIC PATIENTS WITH SYMPTOMATIC HUMAN IMMUNODEFICIENCY VIRUS (HIV) DISEASE

I-48 PEDIATRIC ANGIOCARDIOGRAPHY

I-49 TREATMENT OF TRAVELERS' DIARRHEA DUE TO SUSCEPTIBLE STRAINS OF ENTEROTOXIGENIC ESCHERICHIA COLI

I-50 FOR USE IN WOMEN WITH AXILLARY NODE-NEGATIVE BREAST CANCER

I-51 TREATMENT OF PRIMARY DYSMENORRHEA AND FOR THE TREATMENT OF IDIOPATHIC HEAVY MENSTRUAL BLOOD LOSS

I-52 PEDIATRIC EXCRETORY UROGRAPHY

I-53 TREATMENT OF PANIC DISORDER, WITH OR WITHOUT AGORAPHOBIA

I-54 RENAL CONCENTRATION CAPACITY TEST

I-55 HYPERTENSION

I-56 EROSION GASTROESOPHAGEAL REFLUX DISEASE

I-57 SHORT-TERM TREATMENT OF ACTIVE DUODENAL ULCER

I-58 INITIAL TREATMENT OF ADVANCED OVARIAN CARCINOMA IN COMBINATION WITH OTHER APPROVED CHEMOTHERAPEUTIC AGENTS

I-59 ENDOSCOPICALLY DIAGNOSED ESOPHAGITIS, INCLUDING EROSION AND ULCERATIVE ESOPHAGITIS, AND ASSOCIATED HEARTBURN DUE TO GASTROESOPHAGEAL REFLUX DISEASE

I-60 SINGLE APPLICATION TREATMENT OF HEAD LICE IN CHILDREN TWO MONTHS TO TWO YEARS IN AGE

I-61 FEMALE ANDROGENETIC ALOPECIA

I-62 PREVENTION AND TREATMENT OF POSTMENOPAUSAL OSTEOPOROSIS

I-63 ONCE DAILY TREATMENT AS INITIAL THERAPY IN THE TREATMENT OF HYPERTENSION

I-64 PREVENTION OF SUPRAVENTRICULAR TACHYCARDIAS

I-65 PREVENTION OF UPPER GASTROINTESTINAL BLEEDING IN CRITICALLY ILL PATIENTS

I-66 UNCOMPLICATED GONORRHEA

I-67 TREATMENT OF ACUTE ASTHMATIC ATTACKS IN CHILDREN SIX YEARS OF AGE AND OLDER

I-68 CENTRAL PRECOCIOUS PUBERTY

I-69 SHORT TERM TREATMENT OF PATIENTS WITH SYMPTOMS OF GASTROESOPHAGEAL REFLUX DISEASE (GERD), AND FOR THE SHORT TERM TREATMENT OF ESOPHAGITIS DUE TO GERD INCLUDING ULCERATIVE DISEASE DIAGNOSED BY ENDOSCOPY

I-70 USE IN COMBINATION WITH 5-FLUOROURACIL TO PROLONG SURVIVAL IN THE PALLIATIVE TREATMENT OF PATIENTS WITH ADVANCED COLORECTAL CANCER

I-71 VARICELLA INFECTIONS (CHICKENPOX)

I-72 PREVENTION OF CMV DISEASE IN TRANSPLANT PATIENTS AT RISK FOR CMV DISEASE

I-73 INITIATE AND MAINTAIN MONITORED ANESTHESIA CARE (MAC) SEDATION DURING DIAGNOSTIC PROCEDURES

I-74 INTRAVENOUS DIGITAL SUBTRACTION ANGIOGRAPHY

I-75 TREATMENT OF ENDOSCOPICALLY DIAGNOSED EROSION ESOPHAGITIS

I-76 PREVENTION OF OSTEOPOROSIS

I-77 DERMAL INFECTIONS-TINEA PEDIS, TINEA CORPORIS, TINEA CRURIS DUE TO EPIDERMOPHYTON FLOCCOSUM

I-78 CONTRAST ENHANCED COMPUTED TOMOGRAPHIC IMAGING OF THE HEAD AND BODY AND INTRAVENOUS EXCRETORY UROGRAPHY

I-79 MANAGEMENT OF CHRONIC STABLE ANGINA AND ANGINA DUE TO CORONARY ARTERY SPASM

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

I-80 DIAGNOSIS AND LOCALIZATION OF ISCHEMIA AND CORONARY HEART DISEASE

I-81 PROPHYLAXIS IN DESIGNATED IMMUNOCOMPROMISED CONDITIONS TO REDUCE THE INCIDENCE OF OROPHARYNGEAL CANDIDIASIS

I-82 TREATMENT OF TRAVELERS' DIARRHEA

I-83 ANGIOCARDIOGRAPHY, CONTRAST ENHANCED COMPUTED TOMOGRAPHIC IMAGING OF THE HEAD AND BODY, AND INTRAVENOUS EXCRETORY UROGRAPHY IN CHILDREN

I-84 INTRAOPERATIVE AND POSTOPERATIVE TACHYCARDIA AND/OR HYPERTENSION

I-85 TREATMENT OF ANOREXIA ASSOCIATED WITH WEIGHT LOSS IN PATIENTS WITH AIDS

I-86 TREATMENT OF SECONDARY CARNITINE DEFICIENCY

I-87 RENAL IMAGING AGENT FOR USE IN CHILDREN

I-88 MANAGEMENT OF ENDOMETRIOSIS

I-89 EPIDURAL USE IN LABOR AND DELIVERY AS AN ANALGESIC ADJUNCT TO BUPIVACAINE

I-90 INTENSIVE CARE UNIT SEDATION

I-91 MONOTHERAPY USE FOR HYPERTENSION

I-92 ADJUNCTIVE THERAPY IN THE MANAGEMENT OF HEART FAILURE

I-93 PREVENTION OF EXERCISE-INDUCED BRONCHOSPASM IN CHILDREN AGES 4-11 YEARS

I-94 USE WITH MRI IN ADULTS TO PROVIDE CONTRAST ENHANCEMENT AND FACILITATE VISUALIZATION OF LESIONS IN THE BODY [EXCLUDING THE HEART]

I-95 TREATMENT OF LEFT VENTRICULAR DYSFUNCTION FOLLOWING MYOCARDIAL INFARCTION

I-96 TREATMENT OF SYMPTOMATIC BENIGN PROSTATIC HYPERPLASIA

I-97 ORAL OR RECTAL USE IN CHILDREN FOR THE EXAMINATION OF THE GASTROINTESTINAL TRACT

I-98 TREATMENT OF CHILDREN WHO HAVE GROWTH FAILURE ASSOCIATED WITH CHRONIC RENAL INSUFFICIENCY

I-99 PEDIATRIC ANESTHESIA IN CHILDREN 3 YEARS AND OLDER

I-100 TO DECREASE THE INCIDENCE OF CANDIDIASIS IN PATIENTS UNDERGOING BONE MARROW TRANSPLANTATION WHO RECEIVE CYTOTOXIC CHEMOTHERAPY AND/OR RADIATION THERAPY

I-101 TREATMENT OF DIABETIC NEPHROPATHY IN PATIENTS WITH TYPE I INSULIN-DEPENDENT DIABETES MELLITUS AND RETINOPATHY

I-102 TREATMENT OF OBSESSIVE-COMPULSIVE DISORDER

I-103 PROPHYLAXIS AGAINST PNEUMOCYSTIS CARINII PNEUMONIA IN INDIVIDUALS WHO ARE IMMUNOCOMPROMISED AND CONSIDERED TO BE AT RISK OF DEVELOPING PNEUMOCYSTIS CARINII PNEUMONIA

I-104 TREATMENT OF PULMONARY AND EXTRAPULMONARY ASPERGILLOSIS IN PATIENTS WHO ARE INTOLERANT OF OR WHO ARE REFRACTORY TO AMPHOTERICIN B THERAPY

I-105 TREATMENT OF METASTATIC CARCINOMA OF THE BREAST AFTER FAILURE OF FIRST-LINE OR SUBSEQUENT CHEMOTHERAPY

I-106 TREATMENT OF ACROMEGALY

I-107 VAGINAL CANDIDIASIS

I-108 EXPANDED USE-FOR ICU PATIENTS UNDERGOING LONG-TERM INFUSION DURING MECHANICAL VENTILATION

I-109 TYPHOID FEVER

I-110 PREVENTION OF NAUSEA AND VOMITING ASSOCIATED WITH RADIOTHERAPY

I-111 TREATMENT OF PAGET'S DISEASE OF BONE

I-112 MANAGEMENT OF MODERATE TO SEVERE PAIN

I-113 TREATMENT OF PROSTATITIS

I-114 USE IN CHILDREN TO VISUALIZE LESIONS WITH ABNORMAL VASCULARITY IN THE BRAIN (INTRACRANIAL LESIONS), SPINE, AND ASSOCIATED TISSUE

I-115 USE IN MRI IN ADULTS TO VISUALIZE LESIONS IN THE HEAD AND NECK

I-116 MAINTENANCE OF HEALING OF EROSIIVE ESOPHAGITIS

I-117 TO SLOW THE PROGRESSION FO CORONARY ATHEROSCLEROSIS IN PATIENTS WITH CORONARY HEART DISEASE

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

I-118 PREVENTION OF DEEP VEIN THROMBOSIS, WHICH MAY LEAD TO PULMONARY EMBOLISM FOLLOWING KNEE REPLACEMENT SURGERY

I-119 TREATMENT OF ANEMIA CAUSED BY UTERINE LEIOMYOMATA IN WOMEN WHO FAIL IRON THERAPY

I-120 MAINTENANCE THERAPY FOR GASTRIC ULCER PATIENTS AT REDUCED DOSAGE AFTER HEALING ACUTE ULCERS

I-121 EXPANDED PATIENT POPULATION -- USE IN ICU PATIENTS

I-122 PSORIASIS OF THE SCALP

I-123 RELIEF OF MILD TO MODERATE PAIN IN PATIENTS AGED 6 MONTHS AND OLDER

I-124 LEUKOCYTE LABELED SCINTIGRAPHY AS AN ADJUNCT IN THE LOCALIZATION OF INTRA-ABDOMINAL INFECTION AND INFLAMMATORY BOWEL DISEASE

I-125 EXPANSION OF CONSCIOUS SEDATION INDICATION TO INCLUDE SHORT THERAPEUTIC PROCEDURES

I-126 ADJUNCT TO THALLIUM- 201 MYOCARDIAL PERFUSION IN PATIENTS UNABLE TO EXERCISE ADEQUATELY

I-127 TREATMENT OF ACYCLOVIR-RESISTANT HERPES IN IMMUNOCOMPROMISED PATIENTS

I-128 IN PT W/ CH DISEASE AND HYPERCHOLESTEROLEMIA: REDUCE RISK TOTAL MORTALITY BY REDUCING CORONARY DEATH; REDUCE RISK NON-FATAL MI; REDUCE RISK UNDERGOING MYOCARDIAL REVASCULARIZATION PROCEDURES; REDUCTION ELEVATED TOTAL AND LDL CHOL LEVELS...

I-129 TREATMENT OF ALCOHOL DEPENDENCE

I-130 MAINTENANCE OF HEALING OF EROSIIVE ESOPHAGITIS

I-131 PERIPHERAL ARTERIOGRAPHY

I-132 TREATMENT OF MANIC PHASE OF BIPOLAR DISORDER

I-133 MANAGEMENT OF CHRONIC STABLE ANGINA

I-134 HEART FAILURE POST MYOCARDIAL INFARCTION

I-135 BONE METASTASES ASSOCIATED WITH MULTIPLE MYELOMA

I-136 IDIOPATHIC CHRONIC URTICARIA

I-137 PREVENTION OF METAL-INDUCED HEART BURN, ACID INDIGESTION, AND SOUR STOMACH WHEN TAKEN 30 MINUTES PRIOR TO CONSUMING FOOD OR BEVERAGES

I-138 TREATMENT OF ACUTE RECURRENT GENITAL HERPES

I-139 PALLIATIVE TREATMENT OF ADVANCED BREAST CANCER IN PRE- AND PERIMENOPAUSAL WOMEN

I-140 PREVENTION OF CYTOMEGALOVIRUS (CMV) DISEASE IN INDIVIDUALS WITH HIV INFECTION AT RISK FOR DEVELOPING CMV DISEASE

I-141 TREATMENT OF HEMODYNAMICALLY STABLE PATIENTS WITHIN 24 HOURS OF ACUTE MYOCARDIAL INFARCTION TO IMPROVE SURVIVAL

I-142 LOCALIZE MYOCARDIAL ISCHEMIA (REVERSIBLE DEFECT) AND INFARCTION (NON-REVERSIBLE DEFECTS) IN EVALUATING MYOCARDIAL FUNCTION

I-143 EPISODIC TREATMENT OF RECURRENT GENITAL HERPES IN IMMUNOCOMPETENT ADULTS

I-144 ENHANCEMENT OF MRI OF THE ADULT BODY INTERNAL ORGANS

I-145 0.1MMOL/KG AS A SINGLE INTRAVENOUS BOLUS FOR MRI OF THE CNS IN CHILDREN

I-146 CONTRAST ENHANCEMENT AND FACILITATION OF VISUALIZATION OF EXTRACRANIAL HEAD AND NECK LESIONS

I-147 PREVENTION OF GALLSTONE FORMATION IN OBESE PATIENTS EXPERIENCING RAPID WEIGHT LOSS

I-148 TREATMENT OF ACUTE PNEUMOCYSTIS CARINI PNEUMONIA (PCP) IN HIV-INFECTED PATIENTS WHOSE ALVEOLAR-ARTERIAL OXYGEN DIFFERENCE (AaDO₂) IS LESS THAN OR EQUAL TO 55 TORR

I-149 TREATMENT OF PATIENTS WITH NON-SMALL CELL LUNG CANCER

I-150 TREATMENT OF OBSESSIVE COMPULSIVE DISORDER AND PANIC DISORDER

I-151 PREVENTION OF AND PREVENTION OF FURTHER POSTOPERATIVE NAUSEA AND VOMITING IN PEDIATRIC PATIENTS RECEIVING GENERAL ANESTHESIA

I-152 SLOWING THE PROGRESSION OF CORONARY ATHEROSCLEROSIS AND REDUCING THE RISK OF ACUTE CORONARY EVENTS

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

I-153 MANAGEMENT OF SEVERE SPASTICITY [ENCOMPASSES SPINAL AND CEREBRAL ORIGIN]

I-154 PATIENT POPULATION ALTERED TO INCLUDE PEDIATRIC USE

I-155 TREATMENT OF ONYCHOMYCOSIS DUE TO DERMATOPHYTES (TINEA UNGUIUM) OF THE TOENAIL WITH OR WITHOUT FINGERNAIL INVOLVEMENT

I-156 ADDITIONAL DATA REGARDING THE SAFE USE OF NORVASC IN PATIENTS WITH HEART FAILURE

I-157 TREATMENT OF ACUTE UNCOMPLICATED CYSTITIS IN FEMALES

I-158 TREATMENT OF OSTEOLYTIC BONE METASTASES OF BREAST CANCER

I-159 FOR HYPERCHOLESTEROLEMIC PATIENTS WITHOUT CLINICALLY EVIDENT HEART DISEASE REDUCE THE RISK OF MYOCARDIAL INFARCTION, REVASCULARIZATION, AND DEATH DUE TO CARDIOVASCULAR CAUSES WITH NO INCREASE IN DEATH FROM NON-CARDIOVASCULAR CAUSES

I-160 TREATMENT OF BACTERIAL CORNEAL ULCERS

I-161 TREATMENT OF ADULT-ONSET OR CHILDHOOD-ONSET ADULT GROWTH HORMONE DEFICIENCY

I-162 FOR USE IN PATIENTS 6-11 YEARS OF AGE

I-163 TREATMENT OF PHOTOPHOBIA

I-164 CHRONIC BACTERIAL PROSTATITIS

I-165 MANAGEMENT OF ADULTS WITH ACTIVE, CLASSIC AND DEFINITIVE RHEUMATOID ARTHRITIS WHO HAVE HAD INSUFFICIENT THERAPEUTIC RESPONSE TO OR ARE INTOLERANT OF AN ADEQUATE TRIAL OF FULL DOSES OF ONE OR MORE NON-STEROIDAL ANTI-INFLAMMATORY DRUGS

I-166 TREATMENT OF BULIMIA

I-167 COMPLICATED INTRA-ABDOMINAL INFECTIONS (USED IN COMBINATION WITH METRONIDAZOLE) CAUSED BY MIXED AEROBIC/ANAEROBIC PATHOGENS

I-168 MANAGEMENT OF LOCALLY CONFINED STAGE B2-C METASTATIC CARCINOMA OF THE PROSTATE (IN COMBINATION WITH LHRH AGONISTS)

I-169 USE IN COMBINATION WITH CORTICOSTEROIDS AS INITIAL CHEMOTHERAPY FOR THE TREATMENT OF PATIENTS WITH PAIN RELATED TO ADVANCED HORMONE-REFRACTORY PROSTATE CANCER

I-170 PROPHYLACTIC USE DURING HEAD LICE EPIDEMICS

I-171 RELIEF OF SYMPTOMS OF THE COMMON COLD

I-172 TREATMENT OF INITIAL EPISODE OF GENITAL HERPES

I-173 PREOPERATIVELY FOR THE PREVENTION OF INFECTION IN TRANSRECTAL PROSTATE BIOPSY

I-174 PELVIC INFLAMMATORY DISEASE

I-175 TREATMENT OF TINEA CORPORIS AND TINEA CRURIS

I-176 TREATMENT OF POSTOPERATIVE INFLAMMATION IN PATIENTS WHO HAVE UNDERGONE CATARACT EXTRACTION

I-177 TX OF MODERATE ACNE VULGARIS IN FEMALES, GREATER OR EQUAL TO 15YRS OF AGE, WHO HAVE NO KNOWN CONTRAINDICATIONS TO ORAL CONTRACEPTIVE THERAPY, DESIRE CONTRACEPTION, HAVE ACHIEVED MENARCHE AND ARE UNRESPONSIVE TO TOPICAL ANTI-ACNE MEDICATIONS

I-178 TREATMENT OF ONYCHOMYCOSIS OF THE FINGERNAIL WITHOUT CONCOMITANT ONCHOMYCOSIS OF THE TOENAIL WITH A PULSE DOSING REGIMEN

I-179 NOSOCOMIAL PNEUMONIA-MILD TO MODERATE AND SEVERE CAUSED BY HAEMOPHILUS INFLUENZAE OR KLEBSIELLA PNEUMONIAE

I-180 TREATMENT OF PLANTAR TINEA PEDIS (MOCCASIN TYPE)

I-181 TREATMENT OF PATIENTS WITH COMPLEX PARTIAL SEIZURES WITH AND WITHOUT SECONDARY GENERALIZATION

I-182 TREATMENT OF GROWTH FAILURE ASSOCIATED WITH TURNER SYNDROME

I-183 MAINTENANCE THERAPY IN THE MANAGEMENT OF MILD TO MODERATE ASTHMA IN PEDIATRIC PATIENTS AGES 6-11

I-184 TREATMENT OF PANIC DISORDER AT A RECOMMENDED DOSE RANGE OF 1 TO 2MG/DAY (MAXIMUM OF 4MG)

I-185 PREVENTION OF OSTEOPOROSIS IN POSTMENOPAUSAL WOMEN

I-186 TREATMENT OF TINEA (PITYRIASIS) VERSICOLOR CAUSED BY OR PRESUMED TO BE CAUSED BY

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

	PITYROSPORUM ORBICULARE (ALSO KNOWN AS MALASSEZIA FURFUR OR M. ORBICULARE)
I-187	PREVENTION OF FRACTURES IN THE TREATMENT OF POSTMENOPAUSAL OSTEOPOROSIS
I-188	TREATMENT OF ACUTE SINUSITIS AND ACUTE EXACERBATION OF CHRONIC SINUSITIS
I-189	TREATMENT OF ACUTE OTITIS MEDIA IN PEDIATRIC PATIENTS
I-190	PLANAR IMAGING AS A SECOND LINE DIAGNOSTIC DRUG AFTER MAMMOGRAPHY TO ASSIST IN THE EVALUATION OF BREAST LESIONS IN PATIENTS WITH AN ABNORMAL MAMMOGRAM OR A PALPABLE BREAST MASS
I-191	ENDOMETRIAL THINNING AGENT PRIOR TO ENDOMETRIAL ABLATION FOR DYSFUNCTIONAL UTERINE BLEEDING
I-192	THE PREVENTION OF DEEP VEIN THROMBOSIS, WHICH MAY LEAD TO PULMONARY EMBOLISM, IN PATIENTS UNDERGOING ABDOMINAL SURGERY WHO ARE AT RISK FOR THROMBOEMBOLIC COMPLICATIONS AND A NEW DOSAGE REGIMEN, 40MG ONCE DAILY, FOR THIS INDICATION
I-193	TREATMENT OF PANIC DISORDER IN A RECOMMENDED DOSE RANGE OF 50 TO 200MG/DAY
I-194	CONGESTIVE HEART FAILURE
I-195	FOR USE OF LANSOPRAZOLE IN COMBINATION WITH CLARITHROMYCIN AND AMOXICILLIN FOR THE ERADICATION OF HELICOBACTER PYLORI IN PATIENTS WITH ACTIVE DUODENAL ULCER DISEASE OR A ONE-YEAR HISTORY OF DUODENAL ULCER
I-196	ACUTE TREATMENT OF ACTIVE BENIGN GASTRIC ULCER
I-197	MAINTENANCE OF HEALING OF DUODENAL ULCER
I-198	FOR THE USE OF LANSOPRAZOLE IN COMBINATION WITH AMOXICILLIN FOR THE ERADICATION OF HELICOBACTER PYLORI IN PATIENTS WITH ACTIVE DUODENAL ULCER DISEASE OR A ONE-YEAR HISTORY OF A DUODENAL ULCER
I-199	MONOTHERAPY AND COMBINATION THERAPY WITH SULFONYLUREA IN THE TREATMENT OF TYPE II DIABETES
I-200	TREATMENT OF TINEA (PITYRIASIS) VERSICOLOR
I-201	EMPIRICAL THERAPY FOR FEBRILE NEUTROPENIC PATIENTS
I-202	SECOND-LINE TREATMENT OF AIDS-RELATED KAPOSI'S SARCOMA
I-203	MAINTENANCE OF REMISSION OF ULCERATIVE COLITIS
I-204	USE IN PEDIATRIC PATIENTS BETWEEN THE AGES OF 6 AND 11 FOR THE TREATMENT OF THE NASAL SYMPTOMS OF SEASONAL AND PERENNIAL ALLERGIC RHINITIS
I-205	INITIAL ANTICONVULSANT TREATMENT OF STATUS EPILEPTICUS
I-206	TREATMENT OF EDEMA ASSOCIATED WITH CHRONIC RENAL FAILURE
I-207	FOR THE SUPPRESSION OF RECURRENT EPISODES OF GENITAL HERPES IN IMMUNOCOMPETENT ADULTS
I-208	TREATMENT OF OBSESSIVE COMPULSIVE DISORDER IN THE PEDIATRIC POPULATION
I-209	PAROXYSMAL SUPRAVENTRICULAR TACHYCARDIA (PSVT)
I-210	TO SLOW THE PROGRESSION OF CORONARY ATHEROSCLEROSIS IN PATIENTS WITH CORONARY HEART DISEASE AS PART OF A TREATMENT STRATEGY TO LOWER TOTAL AND LDL CHOLESTEROL TO TARGET LEVELS
I-211	FOR USE IN PEDIATRIC POPULATION
I-212	TREATMENT OF SYMPTOMS OF DRY MOUTH IN PATIENTS WITH SJOGREN'S SYNDROME
I-213	TEMPORARY RELIEF OF PAIN AND PHOTOPHOBIA IN PATIENTS UNDERGOING CORNEAL REFRACTIVE SURGERY
I-214	TREATMENT OF OSTEOPOROSIS
I-215	PRE-PROCEDURAL APPLICATION TO ADULT MALE GENITAL SKIN PRIOR TO SITE-SPECIFIC SUBCUTANEOUS INFILTRATION WITH LIDOCAINE FOR THE REMOVAL OF GENITAL WARTS
I-216	FOR THE LONG-TERM TWICE-DAILY (MORNING AND EVENING) ADMINISTRATION IN THE MAINTENANCE TREATMENT OF BRONCHOSPASM ASSOCIATED WITH COPD, INCLUDING CHRONIC BRONCHITIS AND EMPHYSEMA
I-217	PREVENTION (DURING AND FOLLOWING HOSPITALIZATION) OF DEEP VEIN THROMBOSIS, WHICH MAY LEAD TO PULMONARY EMBOLISM, IN PATIENTS UNDERGOING HIP REPLACEMENT SURGERY
I-218	USE OF LIPITOR AS AN ADJUNCTIVE THERAPY TO DIET FOR THE TREATMENT OF PATIENTS WITH ELEVATED SERUM TRIGLYCERIDE LEVELS (FREDERICKSON TYPE IV)

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

I-219	USE OF LIPITOR BY PATIENTS WITH PRIMARY DYSBETALIPOPROTEINEMIA (FREDERICKSON TYPE III) WHO DO NOT RESPOND ADEQUATELY TO DIET
I-220	TREATMENT OF EPISODIC- HEARTBURN, ACID INDIGESTION AND SOUR STOMACH
I-221	TREATMENT OF BENIGN PROSTATIC HYPERPLASIA (BPH) IN MEN WITH AN ENLARGED PROSTATE TO IMPROVE SYMPTOMS, REDUCE THE RISK OF ACUTE URINARY RETENTION AND REDUCE THE RISK OF THE NEED OF SURGERY
I-222	PREVENTION OF ISCHEMIC COMPLICATIONS OF UNSTABLE ANGINA AND NON-Q-WAVE MYOCARDIAL INFARCTION, WHEN CONCURRENTLY ADMINISTERED WITH ASPIRIN
I-223	USE IN THE SYMPTOMATIC RELIEF OF RHINORRHEA ASSOCIATED WITH ALLERGIC AND NONALLERGIC-PERENNIAL RHINITIS IN CHILDREN AGE 6-11 YEARS
I-224	FOR THE USE IN PEDIATRIC PATIENTS 4 TO 11 YEARS OF AGE FOR THE MANAGEMENT OF THE NASAL SYMPTOMS OF SEASONAL AND PERENNIAL ALLERGIC RHINITIS
I-225	USE IN PATIENTS WITH PREVIOUS MI AND NORMAL CHOLESTEROL LEVELS, TO REDUCE RISK OF RECURRENT MI, MYOCARDIAL REVASCULARIZATION, AND CEREBROVASCULAR DISEASE EVENTS
I-226	FIRST-LINE THERAPY FOR THE TREATMENT OF ADVANCED CARCINOMA OF THE OVARY IN COMBINATION WITH CISPLATIN
I-227	SHORT-TERM TREATMENT OF SYMPTOMATIC GASTROESOPHAGEAL REFLUX DISEASE (GERD)
I-228	PREVENTION OF MEAL INDUCED HEARTBURN AT A DOSE OF 75MG TAKEN 30-60MIN PRIOR TO A MEAL
I-229	PRILOSEC (OMEPRAZOLE), AMOXICILLIN, AND CLARITHROMYCIN FOR THE ERADICATION OF H. PYLORI IN PATIENTS WITH DUODENAL ULCER DISEASE
I-230	IN COMBINATION WITH CIS-PLATIN, FOR THE FIRST LINE TREATMENT OF NON-SMALL CELL LUNG CANCER IN PATIENTS WHO ARE NOT CANDIDATES FOR POTENTIALLY CURATIVE SURGERY AND/OR RADIATION
I-231	TREATMENT OF PATIENTS WITH LOCALLY ADVANCED OR METASTATIC BREAST CANCER AFTER FAILURE OF PRIOR CHEMOTHERAPY
I-232	TREATMENT OF RECURRENT MUCOCUTANEOUS HERPES SIMPLEX INFECTIONS IN HIV-AFFECTED PATIENTS AT A DOSE OF 500MG TWICE DAILY
I-233	PROPHYLACTIC USE TO REDUCE PERIOPERATIVE BLOOD LOSS AND THE NEED FOR BLOOD TRANSFUSION IN PATIENTS UNDERGOING CARDIOPULMONARY BYPASS IN THE COURSE OF CORONARY ARTERY BYPASS GRAFT SURGERY
I-234	FOR USE IN COMBINATION WITH CISPLATIN FOR THE FIRST-LINE TREATMENT OF PATIENTS WITH INOPERABLE LOCALLY ADVANCED (STAGE IIIA OR IIIB) OR METASTATIC (STAGE IV) NON-SMALL CELL LUNG CANCER
I-235	PREVENTION OF EXERCISE-INDUCED BRONCHOSPASM IN PATIENTS 12 YEARS OF AGE AND OLDER
I-236	PREVENTION OF EXERCISE-INDUCED BRONCHOSPASM IN PATIENTS 4 YEARS OF AGE AND OLDER
I-237	MAINTENANCE TREATMENT OF ASTHMA AND PREVENTION OF BRONCHOSPASM IN PATIENTS 4 YEARS OF AGE AND OLDER
I-238	ADJUNCTIVE TREATMENT OF LENNOX-GASTAUT SYNDROME IN PEDIATRIC AND ADULT PATIENTS
I-239	TREATMENT OF PATIENTS WITH HOMOZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA
I-240	MANAGEMENT OF SECONDARY HYPERPARATHYROIDISM AND RESULTANT METABOLIC BONE DISEASE IN PATIENTS WITH MODERATE TO SEVERE CHRONIC RENAL FAILURE (CCR 15 TO 55ML/MIN) NOT YET ON DIALYSIS
I-241	USE IN PHOTODYNAMIC THERAPY (PDT) FOR REDUCTION OF OBSTRUCTION AND PALLIATION OF SYMPTOMS IN PATIENTS WITH COMPLETELY OR PARTIALLY OBSTRUCTING ENDOBRONCHIAL NONSMALL CELL LUNG CANCER
I-242	TREATMENT OF MODERATE TO SEVERE VASOMOTOR SYMPTOMS ASSOCIATED WITH THE MENOPAUSE AND IN THE TREATMENT OF VULVAR AND VAGINAL ATROPHY IN WOMEN WITH AN INTACT UTERUS
I-243	USE IN THE SYMPTOMATIC RELIEF OF RHINORRHEA ASSOCIATED WITH THE COMMON COLD IN CHILDREN AGE 5 TO 11 YEARS
I-244	REDUCE THE INCIDENCE OF BREAST CANCER IN WOMEN AT HIGH RISK FOR BREAST CANCER
I-245	TREATMENT OF ACUTE SINUSITIS
I-246	TREATMENT OF UNCOMPLICATED URINARY TRACT INFECTIONS

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

I-247	USE IN CONVERSION TO MONOTHERAPY IN ADULTS WITH PARTIAL SEIZURES WHO ARE RECEIVING TREATMENT WITH A SINGLE ENZYME-INDUCING ANTIEPILEPTIC DRUG
I-248	INPATIENT TREATMENT OF ACUTE DEEP VEIN THROMBOSIS WITH/WITHOUT PULMONARY EMBOLISM WHEN ADMIN WITH WARFARIN SODIUM AND OUTPATIENT TREATMENT OF ACUTE DEEP VEIN THROMBOSIS WITHOUT PULMONARY EMBOLISM WHEN ADMIN WITH WARFARIN SODIUM
I-249	TREATMENT OF CHRONIC HEPATITIS C IN PATIENTS WITH COMPENSATED LIVER DISEASE PREVIOUSLY UNTREATED WITH ALPHA INTERFERON THERAPY
I-250	PRIMARY PREVENTION OF CORONARY HEART DISEASE IN PATIENTS WITHOUT SYMPATOMATIC CARDIOVASCULAR DISEASE WHO HAVE AVERAGE TO MODERATELY ELEVATED TOTAL-C AND LDL-C AND BELOW AVERAGE HDL-C
I-251	TREATMENT OF GENERALIZED ANXIETY DISORDER
I-252	NEW COMBINATION USE OF PRECOSE FOR PATIENTS WITH TYPE 2 DIABETES TREATED WITH DIET PLUS METFORMIN
I-253	COMBINATION USE OF PRECOSE FOR PATIENTS WITH TYPE 2 DIABETES TREATED WITH DIET PLUS INSULIN
I-254	PREVENTION OF POSTMENOPAUSAL OSTEOPOROSIS (LOSS OF BONE MASS)
I-255	PREVENTION OF PNEUMOCYSTIS CARINII PNEUMONIA (PCP)
I-256	USE IN TREATMENT OF SMALL CELL LUNG CANCER SENSITIVE DISEASE AFTER FAILURE OF FIRST-LINE CHEMOTHERAPY
I-257	TREATMENT OF CHRONIC HEPATITIS B ASSOCIATED WITH EVIDENCE OF HEPATITIS B VIRAL REPLICATION AND ACTIVE LIVER INFLAMMATION
I-258	FOR PERENNIAL NONALLERGIC RHINITIS FOR AGES 4 AND ABOVE
I-259	PROPHYLAXIS OF DEEP VEIN THROMBOSIS (DVT), WHICH MAY LEAD TO PULMONARY EMBOLISM, IN PATIENTS UNDERGOING HIP REPLACEMENT SURGERY
I-260	EXPANDED PEDIATRIC USE IN CHILDREN YOUNGER THAN ONE MONTH OF AGE TO BIRTH (WITH A GESTATIONAL AGE OF 37 WEEKS OR GREATER)
I-261	TREATMENT OF SOCIAL ANXIETY DISORDER
I-262	TREATMENT OR PREVENTION OF BRONCHOSPASM WITH REVERSIBLE OBSTRUCTIVE AIRWAY DISEASE AND FOR THE PREVENTION OF EXERCISE INDUCED BRONCHOSPASM IN CHILDREN AGES 4-12
I-263	TREATMENT OF UNSTABLE ANGINA AND NON-Q-WAVE MYOCARDIAL INFARCTION FOR THE PREVENTION OF ISCHEMIC COMPLICATIONS IN PATIENTS ON CONCURRENT ASPIRIN THERAPY
I-264	PREVENTION OF NAUSEA AND VOMITING ASSOCIATED WITH RADIATION, INCLUDING TOTAL BODY IRRADIATION (TBI) AND FRACTIONATED ABDOMINAL RADIATION
I-265	TREATMENT OF ATOPIC DERMATITIS IN PEDIATRIC PATIENTS 6 YEARS AND OLDER
I-266	USE OF TOPAMAX AS ADJUNCTIVE THERAPY IN PEDIATRIC PATIENTS AGES 2-16 YEARS WITH PARTIAL ONSET SEIZURES
I-267	USE IN PEDIATRIC PATIENTS 3 MONTHS OLD AND OLDER - FOR CORTICOSTEROID-RESPONSIVE DERMATOSES
I-268	PROPHYLAXIS AND CHRONIC TREATMENT OF ASTHMA IN PATIENTS 7-11 YEARS OF AGE
I-269	PREVENTION OF NAUSEA AND VOMITING ASSOCIATED WITH HIGHLY EMETOGENIC CANCER CHEMOTHERAPY, INCLUDING CISPLATIN
I-270	ADJUVANT TREATMENT OF NODE-POSITIVE BREAST CANCER ADMINISTRED SEQUENTIALLY TO STANDARD DOXORUBICIN-CONTAINING COMBINATION CHEMOTHERAPY
I-271	TREATMENT OF OSTEOPOROSIS IN POSTMENOPAUSAL WOMEN
I-272	TREATMENT OF GLUCOCORTICOID-INDUCED OSTEOPOROSIS IN MEN AND WOMEN RECEIVING GLUCOCORTICOID IN A DAILY DOSE EQUIVALENT TO 7.5MG OR GREATER OF PREDNISONE AND WHO HAVE LOW BONE MINERAL DENSITY
I-273	ADJUNCT TO DIET TO INCREASE HDL-C IN PATIENTS WITH PRIMARY HYPERCHOLESTEROLEMIA (HETEROZYGOUS FAMILIAL AND NON FAMILIAL) AND MIXED DYSLIPIDEMIA (FREDERICKSON TYPES IIA AND IIB)
I-274	USE OF TOPAMAX AS ADJUNCTIVE THERAPY IN THE TREATMENT OF PRIMARY GENERALIZED TONIC-CLONIC SEIZURES
I-275	USE IN COMBINATION WITH METFORMIN AND SULFONYLUREA IN PATIENTS WITH TYPE 2 DIABETES
I-276	USE OF REZULIN IN COMBINATION WITH METFORMIN AND SULFONYLUREAS IN PATIENTS WITH

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

TYPE 2 DIABETES

- I-277 TREATMENT OF TYPE III HYPERLIPOPROTEINEMIA
- I-278 TREATMENT OF PATIENTS WITH ISOLATED HYPERTRIGLYCERIDEMIA (FREDERICKSON TYPE IV)
- I-279 TREATMENT OF POST-TRAUMATIC STRESS DISORDER
- I-280 USE OF CARNITOR INJECTION FOR THE PREVENTION AND TREATMENT OF CARNITINE DEFICIENCY IN PATIENTS WITH END STAGE RENAL DISEASE WHO ARE UNDERGOING DIALYSIS
- I-281 INCREASING HDL-C IN PATIENTS WITH PRIMARY HYPERCHOLESTEROLEMIA (HETEROZYGOUS FAMILIAL AND NONFAMILIAL) AND MIXED DYSLIPIDEMIA (FREDERICKSON TYPES IIA AND IIB)
- I-282 TREATMENT OF PATIENTS WITH LOCALLY ADVANCED OR METASTATIC NON-SMALL CELL LUNG CANCER AFTER FAILURE OF PRIOR PLATINUM-BASED CHEMOTHERAPY
- I-283 TO REDUCE THE INCIDENCE OF MODERATE TO SEVERE XEROSTOMIA IN PATIENTS UNDERGOING POST-OPERATIVE RADIATION TREATMENT FOR HEAD AND NECK CANCER, WHERE THE RADIATION PORT INCLUDES A SUBSTANTIAL PORTION OF THE PAROTID GLANDS
- I-284 TO REDUCE THE NUMBER OF ADENOMATOUS COLORECTAL POLYPS IN FAMILIAL ADENOMATOUS POLYPOSIS PATIENTS AS AN ADJUNCT TO USUAL CARE
- I-285 TREATMENT OF NASAL SYMPTOMS OF SEASONAL AND PERENNIAL RHINITIS IN ADULTS AND CHILDREN 3 YEARS OF AGE AND OLDER
- I-286 TREATMENT OF PATIENTS WITH FREDERICKSON TYPE III
- I-287 USE OF PRAVASTATIN IN PATIENTS WITH EVIDENT CORONARY HEART DISEASE TO REDUCE THE RISK OF TOTAL MORTALITY BY REDUCING CORONARY DEATH
- I-288 CHANGES IN SEVERAL SECTIONS OF THE INSERT TO INCORPORATE STATEMENTS CONCERNING THE USE OF HIGH DOSES OF LISINAPRIL TO REDUCE THE RISK OF THE COMBINED OUTCOMES OF MORTALITY AND HOSPITALIZATION IN PATIENTS WITH CONGESTIVE HEART FAILURE
- I-289 USE OF AVANDIA IN COMBINATION WITH A SULFONYLUREA IN PATIENTS WITH TYPE 2 DIABETES MELLITUS WHEN DIET AND EXERCISE WITH EITHER SINGLE AGENT DOES NOT ACHIEVE ADEQUATE GLYCEMIC CONTROL
- I-290 PREVENTION OF CORTICOSTEROID-INDUCED OSTEOPOROSIS
- I-291 PREVENTION OF POSTMENOPAUSAL OSTEOPOROSIS
- I-292 TREATMENT OF POSTMENOPAUSAL OSTEOPOROSIS
- I-293 TREATMENT OF CORTICOSTEROID-INDUCED OSTEOPOROSIS
- I-294 TREATMENT OF UNCOMPLICATED ACUTE ILLNESS DUE TO INFLUENZA A AND B IN PEDIATRIC PATIENTS 7 YEARS AND OLDER WHO HAVE BEEN SYMPTOMATIC FOR NO MORE THAN 2 DAYS
- I-295 PREVENTION OF POSTMENOPAUSAL OSTEOPOROSIS FOR WOMEN WITH AN INTACT UTERUS
- I-296 LONG-TERM INTRAVENOUS TREATMENT OF PULMONARY HYPERTENSION ASSOCIATED WITH THE SCLERODERMA SPECTRUM OF DISEASE IN NYHA CLASS III AND CLASS IV PATIENTS WHO DO NOT RESPOND TO CONVENTIONAL THERAPY
- I-297 SHORT-TERM TREATMENT OF ACUTE MANIC EPISODES ASSOCIATED WITH BIPOLAR I DISORDER
- I-298 TREATMENT OF PATIENTS WITH FREDERICKSON TYPE IIA AND IIB HYPERLIPOPROTEINEMIA
- I-299 USE OF CAMPTOSAR AS A COMPONENT OF FIRST-LINE THERAPY IN COMBINATION WITH 5-FLUOROURACIL AND LEUCOVORIN FOR PATIENTS WITH METASTATIC CARCINOMA OF THE COLON OR RECTUM
- I-300 PROPHYLAXIS FOR ASTHMA IN CHILDREN 2-5 YEARS OF AGE
- I-301 TREATMENT OF SIGNS AND SYMPTOMS OF ALLERGIC CONJUNCTIVITIS
- I-302 TREATMENT OF PEDIATRIC PATIENTS WITH PRADER-WILLI SYNDROME
- I-303 INCREASING HDL-CHOLESTEROL IN PATIENTS WITH PRIMARY HYPERCHOLESTEROLEMIA AND MIXED DYSLIPIDEMIAS
- I-304 TREATMENT OF PATIENTS WITH FREDERICKSON TYPE IV
- I-305 TREATMENT OF LEVOFLOXACIN SUSCEPTIBLE STRAINS OF PENICILLIN-RESISTANT STREPTOCOCCUS PNEUMONIAE IN PATIENTS WITH COMMUNITY ACQUIRED PNEUMONIA
- I-306 INDUCTION OF SPERMATOGENESIS IN MEN WITH PRIMARY AND SECONDARY HYPOGONADOTROPIC HYPOGONADISM IN WHOM THE CAUSE OF INFERTILITY IS NOT DUE TO PRIMARY TESTICULAR FAILURE
- I-307 NEW COMBINATION USE OF METFORMIN AND INSULIN IN TYPE 2 DIABETES

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

I-308 TREATMENT OF PEDIATRIC PATIENTS WITH POLYARTICULAR COURSE JUVENILE RHEUMATOID ARTHRITIS WHO RESPONDED INADEQUATELY TO SALICYLATES OR OTHER NSAIDS

I-309 USE OF ACTONEL 35MG ONCE A WEEK TO INCREASE BONE MASS IN MEN WITH OSTEOPOROSIS

I-310 REDUCTION IN RISK OF MYOCARDIAL INFARCTION, STROKE, AND DEATH FROM CARDIOVASCULAR CAUSES

I-311 ADJUNCTIVE THERAPY IN THE TREATMENT OF PARTIAL SEIZURES IN PEDIATRIC PATIENTS AGE 3 TO 12 YEARS

I-312 FIRST LINE TREATMENT OF POSTMENOPAUSAL WOMEN WITH HORMONE RECEPTOR POSITIVE OR HORMONE RECEPTOR UNKNOWN LOCALLY ADVANCED OR METASTATIC BREAST CANCER

I-313 EXTENSION OF INDICATION TO PROVIDE FOR MAINTENANCE OF RESPONSE

I-314 TOPICAL ANESTHETIC FOR SUPERFICIAL MINOR SURGERY OF GENITAL MUCOUS MEMBRANES AND AS AN ADJUNCT FOR LOCAL INFILTRATION ANESTHESIA IN GENITAL MUCOUS MEMBRANES

I-315 THROMBOPROPHYLAXIS OF DEEP VEIN THROMBOSIS, WHICH MAY LEAD TO PULMONARY EMBOLISM, IN MEDICAL PATIENTS WHO ARE AT RISK FOR THROMBOEMBOLIC COMPLICATIONS DUE TO SEVERELY RESTRICTED MOBILITY DURING ACUTE ILLNESS

I-316 TREATMENT OF NSAID-ASSOCIATED GASTRIC ULCER PATIENTS WHO CONTINUE NSAID USE AND REDUCING RISK OF NSAID-ASSOCIATED GASTRIC ULCERS IN PATIENTS WITH HISTORY OF DOCUMENTED GASTRIC ULCER WHO REQUIRE USE OF AN NSAID

I-317 PROPHYLAXIS OF INFLUENZA IN ADULTS AND ADOLESCENTS 13 YEARS AND OLDER

I-318 FIRSTLINE TREATMENT OF POSTMENOPAUSAL WOMEN WITH HORMONE RECEPTOR POSITIVE OR HORMONE RECEPTOR UNKNOWN LOCALLY ADVANCED OR METASTATIC BREAST CANCER

I-319 USE FOR SUSPECTED OR CONFIRMED METHANOL POISONING, EITHER ALONE OR IN COMBINATION WITH HEMODIALYSIS

I-320 TREATMENT OF TYPE 2 DIABETES IN PEDIATRIC PATIENTS (AGES 10-16 YEARS)

I-321 JUVENILE RHEUMATOID ARTHRITIS

I-322 USE OF DIPRIVAN IN PATIENTS 3 MONTHS TO 16 YEARS

I-323 COLORECTAL CANCER

I-324 REDUCING NEUROLOGIC DISABILITY AND/OR FREQUENCY OF CLINICAL RELAPSES IN PATIENTS WITH SECONDARY (CHRONIC) PROGRESSIVE, PROGRESSIVE RELAPSING, OR WORSENING RELAPSING-REMITTING MULTIPLE SCLEROSIS

I-325 PREVENTION OF RELAPSE AND RECURRENCE OF DEPRESSION

I-326 GENERALIZED ANXIETY DISORDER

I-327 SYMPTOMATIC RELIEF OF RHINORRHEA ASSOCIATED WITH SEASONAL ALLERGIC RHINITIS IN PATIENTS 5 YEARS AND OLDER

I-328 PROPHYLAXIS AND CHRONIC TREATMENT OF ASTHMA IN PATIENTS 5-6 YEARS OF AGE

I-329 UNCOMPLICATED SKIN AND SKIN STRUCTURE INFECTIONS

I-330 MAINTENANCE OF HEALING OF EROSIIVE ESOPHAGITIS AND CONTROL OF DAYTIME AND NIGHTTIME HEARTBURN SYMPTOMS IN PATIENTS WITH GERD

I-331 TREATMENT OF MODERATE ACNE VULGARIS

I-332 EMPIRIC THERAPY IN FEBRILE NEUTROPENIC PATIENTS WITH SUSPECTED FUNGAL INFECTIONS (EFTN)

I-333 TOPICAL TREATMENT OF TINEA (PITYRIASIS) VERSICOLOR DUE TO MALASSEZIA FURFUR (FORMERLY PITYROSPORUM ORBICULARE)

I-334 LONG-TERM TREATMENT OF GROWTH FAILURE IN CHILDREN BORN SMALL FOR GESTATIONAL AGE WHO FAIL TO MANIFEST CATCH-UP GROWTH BY TWO YEARS OF AGE

I-335 ADJUNCTIVE THERAPY IN PATIENTS TWO YEARS AND OLDER WITH SEIZURES ASSOCIATED WITH LENNOX-GASTAUT SYNDROME

I-336 EXPANSION OF INDICATION TO INCLUDE THE TREATMENT OF PATIENTS WITH PREDOMINATELY CLASSIC SUBFOVEAL CHOROIDAL NEOVASCULARIZATION DUE TO PATHOLOGIC MYOPIA OR PRESUMED OCULAR HISTOPLASMOSIS

I-337 PATHOLOGICAL HYPERSECRETION ASSOCIATED WITH ZOLLINGER-ELLISON SYNDROME

I-338 MANAGEMENT OF ACUTE PAIN IN ADULTS AND TREATMENT OF PRIMARY DYSMENORRHEA

I-339 TREATMENT OF HEPATITIS B IN PEDIATRIC PATIENTS AGES 2-17 YEARS

I-340 ATOPIC DERMATITIS IN PEDIATRIC PATIENTS AGES 2-5

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

I-341 BREAST CANCER COMBINATION THERAPY

I-342 USE OF FORADIL FOR LONG-TERM, TWICE DAILY (MORNING AND EVENING) ADMINISTRATION IN THE MAINTENANCE TREATMENT OF BRONCHO-CONSTRICTION IN PATIENTS WITH COPD INCLUDING CHRONIC BRONCHITIS AND EMPHYSEMA

I-343 USE OF COREG FOR SEVERE HEART FAILURE

I-344 ACNE VULGARIS

I-345 TREATMENT OF POSTTRAUMATIC STRESS DISORDER

I-346 TREATMENT OF SYMPTOMATIC GASTRO ESOPHAGEAL REFLUX DISEASE (GERD)

I-347 TREATMENT OR PREVENTION OF BRONCHOSPASM IN CHILDREN 6 YEARS OF AGE AND OLDER WITH OBSTRUCTIVE AIRWAY DISEASE

I-348 LONG-TERM, TWICE-DAILY (MORNING AND EVENING) ADMINISTRATION IN THE MAINTENANCE TREATMENT OF BRONCHOSPASM ASSOCIATED WITH COPD (INCLUDING EMPHYSEMA AND CHRONIC BRONCHITIS)

I-349 ACUTE CORONARY SYNDROME

I-350 TREATMENT OF HETEROZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA IN ADOLESCENT BOYS AND GIRLS AT LEAST ONE YEAR POSTMENARCHAL, AGES 10 TO 17 YEARS, WITH A RECOMMENDED DOSING RANGE OF 10 TO 40MG ONCE DAILY

I-351 PREVENTION OF POSTMENOPAUSAL OSTEOPOROSIS FOR ALL STRENGTHS

I-352 ANTICOAGULANT IN PATIENTS WITH OR AT RISK FOR HEPARIN-INDUCED THROMBOCYTOPENIA UNDERGOING PERCUTANEOUS CORONARY INTERVENTIONS (PCI)

I-353 TREATMENT OF SIGNS AND SYMPTOMS OF RHEUMATOID ARTHRITIS

I-354 MANAGEMENT OF POST HERPETIC NEURALGIA

I-355 PREMENSTRUAL DYSPHORIC DISORDER

I-356 TREATMENT OF PATHOLOGICAL HYPERSECRETORY CONDITIONS, INCLUDING ZOLLINGER-ELLISON SYNDROME

I-357 TREATMENT OF COMPLICATED SKIN AND SKIN STRUCTURE INFECTIONS

I-358 TREATMENT OF PANIC DISORDER

I-359 TREATMENT OF VULVAR AND VAGINAL ATROPHY ASSOCIATED WITH MENOPAUSE

I-360 TREATMENT OF NASAL SYMPTOMS OF SEASONAL AND PERENNIAL RHINITIS IN CHILDREN AGES TWO UP TO AGE THREE

I-361 TREATMENT OF MULTIPLE MYELOMA AND DOCUMENTED BONE METASTASES FROM SOLID TUMORS, IN CONJUNCTION WITH STANDARD ANTINEOPLASTIC THERAPY. PROSTATE CANCER SHOULD HAVE PROGRESSED AFTER TREATMENT WITH AT LEAST ONE HORMONAL THERAPY

I-362 TREATMENT OF PANIC DISORDER, WITH OR WITHOUT AGORAPHOBIA

I-363 ADJUVANT TREATMENT OF POST MENOPAUSAL WOMEN WITH HORMONE RECEPTOR POSITIVE EARLY BREAST CANCER

I-364 TREATMENT OF COMMUNITY-ACQUIRED PNEUMONIA IN ADULTS

I-365 TREATMENT OF HEART FAILURE (NYHA CLASS II-IV) IN PATIENTS WHO ARE INTOLERANT TO AN ACE INHIBITOR

I-366 PREVENTION OF RELAPSE FOLLOWING LONG-TERM TREATMENT OF MAJOR DEPRESSIVE DISORDER

I-367 COMBINATION THERAPY WITH THIAZOLIDINEDIONE TO LOWER BLOOD GLUCOSE IN PTS WHOSE HYPERGLYCEMIA CANNOT BE CONTROLLED BY DIET/EXERCISE PLUS MONOTHERAPY WITH ANY OF THE FOLLOWING AGENTS: METFORMIN, SULFONYLUREAS, REPAGLINIDE, OR THIAZOLIDINEDIONES

I-368 USE OF GLUCOVANCE WITH A THIAZOLIDINEDIONE WHEN GLYCEMIC CONTROL IS NOT OBTAINED WITH GLUCOVANCE ALONE

I-369 PREVENTION AND TREATMENT OF POSTOPERATIVE NAUSEA AND VOMITING

I-370 TREATMENT OF HETEROZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA IN CHILDREN, AGES 8-13 YEARS, WITH RECOMMENDED DOSE OF 20MG ONCE DAILY AND IN ADOLESCENTS, AGES 14-18 WITH A RECOMMENDED DOSE OF 40MG ONCE DAILY

I-371 HELICOBACTER PYLORI ERADICATION TO REDUCE THE RISK OF DUODENAL ULCER RECURRENCE

I-372 NOSOCOMIAL PNEUMONIA

I-373 TREATMENT OF TYPE 2 DIABETIC NEPHROPATHY

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

- I-374 SHORT TERM TOPICAL TREATMENT OF MILD TO MODERATE PLAQUE-TYPE PSORIASIS OF NON SCALP REGIONS
- I-375 FIRST LINE THERAPY FOR THE REDUCTION OF INTRAOCULAR PRESSURE IN PATIENTS WITH OPEN-ANGLE GLAUCOMA OR OCULAR HYPERTENSION
- I-376 TREATMENT OF NEWLY DIAGNOSED ADULT PATIENTS WITH PHILADELPHIA CHROMOSOME POSITIVE CHRONIC MYELOID LEUKEMIA (CML)
- I-377 USE OF BRAVELLE FOR MULTIPLE FOLLICULAR DEVELOPMENT (CONTROLLED OVARIAN STIMULATION) DURING ASSISTED REPRODUCTIVE TECHNOLOGY CYCLES IN PATIENTS WHO HAVE PREVIOUSLY RECEIVED PITUITARY SUPPRESSION
- I-378 RELIEF OF SYMPTOMS OF SEASONAL ALLERGIC RHINITIS IN ADULTS AND PEDIATRIC PATIENTS 2 YEARS OF AGE AND OLDER
- I-379 USE TAXOTERE IN COMBINATION WITH CISPLATIN FOR THE TREATMENT OF PATIENTS WITH UNRESECTABLE, LOCALLY ADVANCED OR METASTATIC NON-SMALL CELL LUNG CANCER WHO HAVE NOT PREVIOUSLY RECEIVED CHEMOTHERAPY FOR THIS CONDITION
- I-380 TO TREAT PATIENTS WITH SCHIZOPHRENIA OR SCHIZOAFFECTIVE DISORDER AT RISK FOR EMERGENT SUICIDAL BEHAVIOR
- I-381 TREATMENT OF COLD SORES (HERPES LABIALIS) IN ADULT AND ADOLESCENT PATIENTS 12 YEARS OF AGE AND OLDER
- I-382 FOR NEWLY-DIAGNOSED HIGH GRADE MALIGNANT GLIOMA PATIENTS AS AN ADJUNCT TO SURGERY AND RADIATION
- I-383 TREATMENT OF TYPE 2 DIABETIC NEPHROPATHY
- I-384 USE IN COMBINATION WITH INSULIN FOR THE TREATMENT OF PATIENTS WITH TYPE 2 DIABETES MELLITUS
- I-385 MODIFICATION OF THE INDICATION FOR COMMUNITY ACQUIRED PNEUMONIA TO ADD "INCLUDING PENICILLIN-RESISTANT STRAINS, MIC PENICILLIN \geq 2MCG/ML" TO STREPTOCOCCUS PNEUMONIAE
- I-386 RAPAMUNE (SIROLIMUS) WITHIN AN IMMUNOSUPPRESSIVE REGIMEN THAT WOULD ALLOW FOR THE WITHDRAWAL OF CYCLOSPORINE 2 TO 4 MONTHS AFTER RENAL TRANSPLANTATION IN PATIENTS CONSIDERED AT LOW TO MODERATE IMMUNOLOGIC RISK FOR RENAL TRANSPLANT REJECTION
- I-387 ADJUNCTIVE THERAPY OF PARTIAL SEIZURES IN PEDIATRIC PATIENTS GREATER THAN OR EQUAL TO 2 YEARS OF AGE
- I-388 TREATMENT OF PATIENTS WITH LEFT VENTRICULAR DYSFUNCTION FOLLOWING MYOCARDIAL INFARCTION
- I-389 SUPPRESSION OF RECURRENT GENITAL HERPES IN HIV-INFECTED INDIVIDUALS
- I-390 USE IN PTS AT HIGH RISK CORONARY EVENTS DUE TO EXISTING CORONARY HEART DISEASE, DIABETES, PERIPHERAL VESSEL DISEASE, STROKE HISTORY, OTHER CV DISEASE TO REDUCE RISK TOTAL MORTALITY BY REDUCING CORONARY DEATH, REDUCE NONFATAL MI & STROKE.....
- I-391 ABLATION OF HIGH-GRADE DYSPLASIA IN BARRETT'S ESOPHAGUS PATIENTS WHO DO NOT UNDERGO ESOPHAGECTOMY
- I-392 TX OF PED PATIENTS W/PH+ CHRONIC PHASE CML DISEASE RECURRENCE AFTER STEM CELL TRANSPLANT OR RESISTANCE TO INTERFERON ALPHA THERAPY. NO CONTROLLED TRIALS DEMONSTRATING A CLINICAL BENEFIT SUCH AS IMPROVEMENT IN DISEASE RELATED SX OR INCREASED SURVIVAL
- I-393 CHRONIC BACTERIAL PROSTATITIS
- I-394 USE IN PATIENTS WITH CORONARY HEART DISEASE TO REDUCE THE RISK OF UNDERGOING CORONARY REVASCULARIZATION PROCEDURES
- I-395 TO IMPROVE PHYSICAL FUNCTION
- I-396 EXPANDED INDICATION TO INCLUDE THE ASSESSMENT OF VENTRICULAR FUNCTION IN SUBJECTS BEING EVALUATED FOR HEART DISEASE AND/OR VENTRICULAR FUNCTION
- I-397 EXTENDED PROPHYLAXIS IN PATIENTS UNDERGOING HIP FRACTURE SURGERY
- I-398 IDIOPATHIC SHORT STATURE
- I-399 TREATMENT OF CANDIDEMIA AND THE FOLLOWING CANDIDA INFECTIONS: INTRA-ABDOMINAL ABSCESES, PERITONITIS AND PLEURAL SPACE INFECTIONS
- I-400 USE OF OLANZAPINE IN COMBINATION WITH LITHIUM OR VALPROATE FOR THE TREATMENT OF ACUTE MANIC EPISODES ASSOCIATED WITH BIPOLAR DISORDER

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

I-401	LONGER-TERM EFFICACY OF ARIPIPIRAZOLE IN THE TREATMENT OF SCHIZOPHRENIA
I-402	DIABETIC FOOT INFECTIONS WITHOUT CONCOMITANT OSTEOMYELITIS
I-403	USE OF VALTREX IN COMBINATION WITH SAFER SEX PRACTICES FOR THE REDUCTION OF THE RISK OF TRANSMISSION OF GENITAL HERPES DURING SUPPRESSIVE THERAPY OF THE SOURCE PARTNER IN A HETEROSEXUAL COUPLE
I-404	MAINTENANCE TREATMENT OF BIPOLAR I DISORDER TO DELAY THE TIME TO OCCURRENCE OF MOOD EPISODES (DEPRESSION, MANIA, HYPOMANIA, MIXED EPISODES) IN PATIENTS TREATED FOR ACUTE MOOD EPISODES WITH STANDARD THERAPY
I-405	TREATMENT OF PREMENSTRUAL DYSPHORIC DISORDER (PMDD) USING AN INTERMITTENT DOSING REGIMEN
I-406	PREVENTION OF CYTOMEGALOVIRUS DISEASE IN KIDNEY, HEART, AND KIDNEY-PANCREAS TRANSPLANT PATIENTS AT HIGH RISK (DONOR CMV SEROPOSITIVE/RECIPIENT CMV SERONEGATIVE)
I-407	IMPROVE SURVIVAL OF STABLE PATIENTS WITH LEFT VENTRICULAR SYSTOLIC DYSFUNCTION (EJECTION FRACTION \leq 40%) AND CLINICAL EVIDENCE OF CONGESTIVE HEART FAILURE AFTER AN ACUTE MYOCARDIAL INFARCTION
I-408	STIMULATION OF PANCREATIC SECRETIONS TO FACILITATE THE IDENTIFICATION OF THE AMPULLA OF VATER AND ACCESSORY PAPILLA DURING ENDOSCOPIC RETROGRADE CHOLANGIO-PANCREATOGRAPHY (ERCP)
I-409	ESOPHAGEAL CANDIDIASIS
I-410	USE OF ADVAIR DISKUS 250/50 FOR CHRONIC OBSTRUCTIVE PULMONARY DISEASE (COPD) ASSOCIATED WITH CHRONIC BRONCHITIS
I-411	EXPANDED INDICATION FOR USE IN COMBINATION WITH ANTIDIABETIC DRUGS IN THE THIAZOLIDINEDIONE CLASS
I-412	MONOTHERAPY FOR THE SHORT TERM TREATMENT OF ACUTE MANIC OR MIXED EPISODES ASSOCIATED WITH BIPOLAR I DISORDER
I-413	ADJUNCTIVE THERAPY FOR THE SHORT TERM TREATMENT OF ACUTE MANIC OR MIXED EPISODES ASSOCIATED WITH BIPOLAR I DISORDER
I-414	PROPHYLAXIS OF DEEP VEIN THROMBOSIS (DVT), WHICH MAY LEAD TO PULMONARY EMBOLISM (PE) IN MEDICAL PATIENTS WHO ARE AT RISK FOR THROMBOEMBOLIC COMPLICATIONS DUE TO SEVERELY RESTRICTED MOBILITY DURING ACUTE ILLNESS
I-415	SEVERE HYPERTENSION WHEN THE VALUE OF ACHIEVING PROMPT BLOOD PRESSURE CONTROL EXCEEDS THE RISK OF INITIATING COMBINATION THERAPY
I-416	THE USE OF CIPRO XR FOR COMPLICATED URINARY TRACT INFECTIONS AND ACUTE UNCOMPLICATED PYELONEPHRITIS
I-417	USE IN THE LONG TERM TREATMENT OF BIPOLAR I DISORDER
I-418	ADJUNCTIVE THERAPY W/ MOOD STABILIZERS (LITHIUM OR DIVALPROEX) IN THE TREATMENT OF ACUTE MANIC EPISODES ASSOCIATED WITH BIPOLAR I DISORDERS
I-419	MONOTHERAPY IN THE TREATMENT OF ACUTE MANIC EPISODES ASSOCIATED WITH BIPOLAR I DISORDER
I-420	TOPICAL TREATMENT OF CLINICALLY TYPICAL, NONHYPERKERATOTIC, NONHYPERTROPHIC ACTINIC KERATOSES ON THE FACE OR SCALP IN IMMUNOCOMPETENT ADULTS
I-421	TREATMENT OF COMPLICATED URINARY TRACT INFECTIONS AND PYELONEPHRITIS DUE TO E.COLI FOR PED PATIENTS (1-17) NOT AS FIRST CHOICE
I-422	INDICATED FOR THE IN-HOSPITAL SHORT-TERM (UP TO 4 HOURS) REDUCTION IN BLOOD PRESSURE IN PEDIATRIC PATIENTS
I-423	ACUTE TREATMENT OF MIGRAINE ATTACKS WITH OR WITHOUT AURA IN ADULTS
I-424	MANAGEMENT OF SECONDARY HYPERPARATHYROIDISM IN PATIENTS WITH MODERATE TO SEVERE CHRONIC RENAL INSUFFICIENCY NOT YET ON DIALYSIS
I-425	ELOXATIN IN COMBINATION WITH INFUSIONAL 5-FLUOROURACIL (5-FU) AND LEUCOVORIN (LV) FOR THE TREATMENT OF PATIENTS PREVIOUSLY UNTREATED FOR ADVANCED COLORECTAL CANCER
I-426	TREATMENT OF ACUTE PULMONARY EMBOLISM WHEN ADMINISTERED IN CONJUNCTION WITH WARFARIN SODIUM
I-427	TREATMENT OF ACUTE DEEP VEIN THROMBOSIS WITHOUT PULMONARY EMBOLISM WHEN ADMINISTERED IN CONJUNCTION WITH WARFARIN SODIUM
I-428	FOR USE IN COMBINATION WITH PACLITAXEL FOR THE FIRST-LINE TREATMENT OF PATIENTS

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

- WITH METASTATIC BREAST CANCER AFTER FAILURE OF PRIOR ANTHRACYCLINE CONTAINING ADJUVANT CHEMOTHERAPY UNLESS ANTHRACYCLINES WERE CLINICALLY CONTRAINDICATED
- I-429 FOR USE IN COMBINATION WITH PREDNISONE FOR THE TREATMENT OF PATIENTS WITH ANDROGEN INDEPENDENT (HORMONE REFRACTORY) METASTATIC PROSTATE CANCER
- I-430 FOR USE IN THE RELIEF OF THE SIGNS AND SYMPTOMS OF RHEUMATOID ARTHRITIS IN ADULTS
- I-431 NOSOCOMIAL PNEUMONIA AND COMMUNITY-ACQUIRED PNEUMONIA CAUSED BY STREPTOCOCCUS PNEUMONIAE INDICATION EXPANDED TO INCLUDE MULTI-DRUG RESISTANT STRAINS
- I-432 TREATMENT OF COMMUNITY ACQUIRED PNEUMONIA CAUSED BY MULTI-DRUG RESISTANT STREPTOCOCCUS PNEUMONIAE
- I-433 TREATMENT OF BIOPSY-CONFIRMED, PRIMARY SUPERFICIAL BASAL CELL CARCINOMA IN IMMUNOCOMPETENT ADULTS, WITH A MAXIMUM TUMOR DIAMETER OF 2.0CM, LOCATED ON THE TRUNK (EXCLUDING ANOGENITAL SKIN), NECK, OR EXTREMITIES (EXCLUDING HANDS AND FEET)
- I-434 PREVENTION OF CARDIOVASCULAR DISEASE IN ADULT PATIENTS WITHOUT CLINICALLY EVIDENT HEART DISEASE, BUT WITH MULTIPLE RISK FACTORS FOR CORONARY HEART DISEASE TO REDUCE RISK OF MI AND RISK FOR REVASCLARIZATION PROCEDURES AND ANGINA
- I-435 CHRONIC IDIOPATHIC CONSTIPATION
- I-436 FOR USE IN COMBINATION WITH DOXORUBICIN AND CYCLOPHOSPHAMIDE FOR THE ADJUVANT TREATMENT OF PATIENTS WITH OPERABLE NODE-POSITIVE BREAST CANCER
- I-437 TREATMENT OF ACUTE MANIC AND MIXED EPISODES ASSOCIATED WITH BIPOLAR DISORDER
- I-438 EMPIRICAL THERAPY FOR PRESUMED FUNGAL INFECTIONS IN FEBRILE, NEUTROPENIC PATIENTS
- I-439 USED TO TREAT ADULTS WITH GROWTH HORMONE DEFICIENCY
- I-440 FOR THE REPLACEMENT OF ENDOGENOUS GROWTH HORMONE IN ADULTS WITH GROWTH HORMONE DEFICIENCY
- I-441 USE COMBINATION WITH INFUSIONAL 5-FU/LV FOR ADJUVANT TREATMENT STAGE III COLON CANCER PTS WHO HAVE UNDERGONE COMPLETE RESECTION PRIMARY TUMOR-BASED ON IMPROVEMENT IN DISEASE FREE SURVIVAL, NO DEMONSTRATED BENEFIT OVERALL SURVIVAL AFTER 4YRS
- I-442 USED FOR CANDIDEMIA IN NONNEUTROPENIC PATIENTS AND THE FOLLOWING CANDIDA INFECTIONS: DISSEMINATED INFECTIONS IN SKIN & INFECTIONS IN ABDOMEN, KIDNEY, BLADDER WALL, AND WOUNDS
- I-443 TREATMENT OF NASAL POLYPS IN PATIENTS 18 YEARS OF AGE AND OLDER
- I-444 USE OF PROTONIX IV FOR INJECTION AS STAND ALONE THERAPY FOR THE SHORT-TERM TREATMENT OF PATIENTS HAVING GASTROESOPHAGEAL REFLUX (GERD) WITH A HISTORY OF EROSIIVE ESOPHAGITIS
- I-445 TO IMPROVE (COMPARED TO 4.25% DEXTROSE) LONG-DWELL ULTRAFILTRATION AND CLEARANCE OF CREATININE AND UREA NITROGEN IN PATIENTS WITH HIGH AVERAGE OR GREATER TRANSPORT CHARACTERISTICS, AS DEFINED USING THE PERITONEAL EQUILIBRATION TEST (PET)
- I-446 EXTENDED ADJUVANT TREATMENT OF EARLY BREAST CANCER IN POSTMENOPAUSAL WOMEN WHO HAVE RECEIVED 5 YRS ADJUVANT TAMOXIFEN THERAPY-EFFECTIVENESS BASED ON AN ANALYSIS OF DISEASE FREE SURVIVAL IN PATIENTS TREATED FOR A MEDIAN 24 MONTHS
- I-447 USE OF COPEGUS (RIBAVIRIN) FOR TREATMENT OF CHRONIC HEPATITIS C IN ADULT PATIENTS COINFECTED WITH HIV IN COMBINATION WITH PEGASYS (PEGINTERFERON ALFA-2A)
- I-448 TREATMENT OF HEART FAILURE (NYHA CLASS II-IV AND EJECTION FRACTION <=40%) TO REDUCE THE RISK OF DEATH FROM CARDIOVASCULAR CAUSES AND TO REDUCE HOSPITALIZATIONS FOR HEART FAILURE
- I-449 TO IMPROVE WAKEFULNESS IN TWO NEW PATIENT POPULATIONS WITH EXCESSIVE SLEEPINESS: THOSE WITH OBSTRUCTIVE SLEEP APNEA/HYPOPNEA SYNDROME AND THOSE WITH SHIFT WORK SLEEP DISORDER
- I-450 TREATMENT OF PATIENTS WITH NEWLY DIAGNOSED HIGH GRADE GLIOMAS CONCOMITANTLY WITH RADIOTHERAPY AND THEN AS ADJUVANT TREATMENT
- I-451 MANAGEMENT OF ENDOMETRIOSIS ASSOCIATED PAIN
- I-452 EXPANDED INDICATION TO INCLUDE TREATMENT OF MULTIPLE MYELOMA PATIENTS WHO HAVE RECEIVED AT LEAST 1 PRIOR THERAPY
- I-453 USE IN COMBINATION WITH A SULFONYLUREA PLUS METFORMIN WHEN DIET, EXERCISE AND

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

- BOTH AGENTS DO NOT RESULT IN ADEQUATE GLYCEMIC CONTROL (TRIPLE THERAPY)
- I-454 MAINTENANCE OF CLINICAL REMISSION OF MILD TO MODERATE CROHN'S DISEASE INVOLVING THE ILEUM AND/OR THE ASCENDING COLON FOR UP TO 3 MONTHS
- I-455 MODIFIED HEART FAILURE INDICATION TO INCLUDE TREATMENT OF HEART FAILURE IN PATIENTS WITH LEFT VENTRICULAR SYSTOLIC DYSFUNCTION (NYHA CLASS II-IV; EJECTION FRACTION LESS THAN OR EQUAL TO 40%)
- I-456 TO REDUCE CARDIOVASCULAR DEATH AND TO REDUCE HEART FAILURE HOSPITALIZATIONS. INCLUDES ADDITIONAL INFORMATION ON THE ADDED EFFECT ON THESE OUTCOMES WHEN USED WITH AN ACE INHIBITOR
- I-457 TREATMENT OF PATIENTS UNDERGOING ABDOMINAL SURGERY WHO ARE AT RISK FOR THROMBOEMBOLIC COMPLICATIONS
- I-458 USE OF BIVALIRUDIN FOR INJECTION WITH PROVISIONAL USE OF GLYCOPROTEIN IIB/IIIA INHIBITOR (GPI) AS LISTED IN THE CLINICAL TRIALS REPLACE-2 SECTION FOR USE AS AN ANTICOAGULANT IN PATIENTS UNDERGOING PERCUTANEOUS CORONARY INTERVENTION (PCI)
- I-459 NON-DIALYSIS DEPENDENT CHRONIC KIDNEY DISEASE (NDD-CKD) PATIENTS RECEIVING OR NOT RECEIVING AN ERYTHROPOIETIN
- I-460 TREATMENT OF DIARRHEA CAUSED BY CRYPTOSPORIDIUM PARVUM IN NON-HIV INFECTED PATIENTS 12 YEARS OF AGE AND OLDER
- I-461 USE AS A SINGLE AGENT FOR ADJUVANT TREATMENT IN PATIENTS WITH DUKES' C COLON CANCER WHO HAVE UNDERGONE COMPLETE RESECTION OF THE PRIMARY TUMOR WHEN TREATMENT WITH FLUOROPYRIMIDINE THERAPY ALONE IS PREFERRED
- I-462 LONG TERM TREATMENT OF IDIOPATHIC SHORT STATURE
- I-463 TREATMENT OF PATIENTS POST MYOCARDIAL INFARCTION
- I-464 TREATMENT OF MODERATE TO SEVERE PRIMARY RESTLESS LEGS SYNDROME
- I-465 PERENNIAL ALLERGIC RHINITIS IN ADULTS AND PEDIATRIC PATIENTS 6 MONTHS OF AGE AND OLDER
- I-466 FOR RELIEF OF THE SIGNS AND SYMPTOMS OF ANKYLOSING SPONDYLITIS
- I-467 USE OF TOPIRAMATE AS INITIAL MONOTHERAPY IN PATIENTS 10 YEARS OF AGE AND OLDER WITH PARTIAL ONSET OR PRIMARY GENERALIZED TONIC CLONIC SEIZURES
- I-468 USE IN PATIENTS WITH STABLE CORONARY ARTERY DISEASE TO REDUCE THE RISK OF CARDIOVASCULAR MORTALITY OR NON-FATAL MYOCARDIAL INFARCTION
- I-469 RELIEF OF THE SIGNS AND SYMPTOMS OF PAUCIARTICULAR OR POLYARTICULAR COURSE JUVENILE RHEUMATOID ARTHRITIS IN PATIENTS 2 YEARS OF AGE AND OLDER
- I-470 DIABETIC PERIPHERAL NEUROPATHIC PAIN
- I-471 INDICATED TO REDUCE THE RISK OF MYOCARDIAL INFARCTION AND STROKE IN PATIENTS WITH TYPE 2 DIABETES AND WITHOUT CLINICALLY EVIDENT CORONARY HEART DISEASE BUT WITH MULTIPLE RISK FACTORS FOR CORONARY HEART DISEASE
- I-472 USE IN PATIENTS WITH ANGIOGRAPHICALLY DOCUMENTED CORONARY ARTERY DISEASE
- I-473 USE IN COMBINATION WITH GEMCITABINE FOR THE FIRST LINE TREATMENT OF PATIENTS WITH LOCALLY ADVANCED UNRESECTABLE OR METASTATIC PANCREATIC CANCER
- I-474 TREATMENT OF IRON DEFICIENCY ANEMIA IN PERITONEAL DIALYSIS DEPENDANT CHRONIC KIDNEY DISEASE IN PATIENTS RECEIVING AN ERYTHROPOIETIN
- I-475 PREVENTION OF NAUSEA AND VOMITING ASSOCIATED WITH INITIAL AND REPEAT COURSES OF MODERATELY EMETOGENIC CANCER CHEMOTHERAPY
- I-476 TREATMENT OF DIABETIC FOOT INFECTIONS WITHOUT OSTEOMYELITIS
- I-477 TREATMENT OF COMPLICATED SKIN AND SKIN STRUCTURE INFECTIONS CAUSED BY METHICILLIN SUSCEPTIBLE STAPHYLOCOCCUS AUREUS, ESCHERICHIA COLI, KLEBSIELLA PNEUMONIAE, OR ENTEROBACTER CLOACAE
- I-478 FOR USE AS ADJUNCTIVE THERAPY IN THE TREATMENT OF PARTIAL SEIZURES IN CHILDREN WITH EPILEPSY AGED 2-4 YEARS
- I-479 TREATMENT OF COMPLICATED INTRA-ABDOMINAL INFECTIONS CAUSED BY E.COLI, B. FRAGILIS, S.ANGINOSUS, S.CONSTELLATUS, E. FAECALIS, P. MIRABILIS, C. PERFRINGENS, B. THETAIOAOMICRON OR PEPTOSTREPTOCOCCUS SPECIES
- I-480 PROPHYLAXIS OF INFLUENZA FOR PATIENTS BETWEEN 1-12 YEARS OF AGE
- I-481 INDICATED FOR THE ADJUVANT TREATMENT OF POSTMENOPAUSAL WOMEN WITH HORMONE RECEPTOR POSITIVE EARLY BREAST CANCER

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

I-482 TREATMENT OF ACUTE MANIC OR MIXED EPISODES ASSOCIATED WITH BIPOLAR I DISORDER WITH OR WITHOUT PSYCHOTIC FEATURES

I-483 PREVENTION OF POSTMENOPAUSAL OSTEOPOROSIS

I-484 FOR THE RISK REDUCTION OF NSAID-ASSOCIATED GASTRIC ULCERS

I-485 TREATMENT OF POSTOPERATIVE INFLAMMATION AND REDUCTION OF OCULAR PAIN IN PATIENTS WHO HAVE UNDERGONE CATARACT EXTRACTION

I-486 ANGIOMAX IS INDICATED FOR PATIENTS WITH, OR AT RISK OF, HIT/HITTS UNDERGOING PCI

I-487 INDICATED FOR THE RELIEF OF THE INFLAMMATORY AND PRURITIC MANIFESTATIONS OF CORTICOSTEROID RESPONSIVE DERMATOSES IN PATIENTS 12 YRS OF AGE OR OLDER

I-488 MAINTENANCE THERAPY IN BIPOLAR I DISORDER

I-489 FOR USE IN PEDIATRIC PATIENTS WITH TYPE I DIABETES

I-490 FOR USE IN COMBINATION WITH CISPLATIN AND FLUOROURACIL FOR THE TREATMENT OF PATIENTS WITH ADVANCED GASTRIC ADENOCARCINOMA, INCLUDING ADENOCARCINOMA OF GASTROESOPHAGEAL JUNCTION, WHO HAVE NOT RECEIVED PRIOR CHEMOTHERAPY FOR ADVANCED DISEASE

I-491 INFLUENZA PROPHYLAXIS

I-492 MONOTHERAPY IN THE TREATMENT OF ACUTE MANIC OR MIXED EPISODES IN BIPOLAR I DISORDER, WITH OR WITHOUT PSYCHOTIC FEATURES

I-493 ADMINISTERED IN COMBINATION WITH FENOFIBRATE, AS ADJUNCTIVE THERAPY TO DIET FOR THE REDUCTION OF ELEVATED TOTAL-C, LDL-C, APO B, AND NON-HDL-C IN PATIENTS WITH MIXED HYPERLIPIDEMIA

I-494 CLINICAL DATA IN SUPPORT OF AVANDAMET AS AN ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN PATIENTS WITH TYPE 2 DIABETES MELLITUS WHEN TREATMENT WITH DUAL ROSIGLITAZONE AND METFORMIN THERAPY IS APPROPRIATE

I-495 ADJUVANT TX OF POSTMENOPAUSAL WOMEN WITH ESTROGEN-RECEPTOR POSITIVE EARLY BREAST CANCER WHO HAVE RECEIVED 2 TO 3 YRS OF TAMOXIFEN AND ARE SWITCHED TO AROMASIN FOR COMPLETION OF A TOTAL OF 5 CONSECUTIVE YRS OF ADJUVANT HORMONAL THERAPY

I-496 LONG TERM TREATMENT OF GROWTH FAILURE ASSOCIATED WITH TURNER SYNDROME IN PATIENTS WHO HAVE OPEN EPIPHYSES

I-497 PREVENTION OF SEASONAL MAJOR DEPRESSIVE EPISODES IN PATIENTS WITH SEASONAL AFFECTIVE DISORDER

I-498 PREVENTION OF POSTOPERATIVE NAUSEA AND VOMITING

I-499 USE OF GEMZAR IN COMBINATION WITH CARBOPLATIN FOR THE TREATMENT OF PATIENTS WITH ADVANCED OVARIAN CANCER THAT HAS RELAPSED AT LEAST 6 MONTHS AFTER COMPLETION OF PLATINUM-BASED THERAPY

I-500 FOR USE IN COMBINATION WITH DEXAMETHASONE FOR THE TREATMENT OF MULTIPLE MYELOMA PATIENTS WHO HAVE RECEIVED AT LEAST ONE PRIOR THERAPY

I-501 TREATMENT OF RECURRENT HERPES LABIALIS (COLD SORES) IN IMMUNOCOMPETENT PATIENTS WITH A SINGLE DOSE OF FAMCICLOVIR 1500 MG.

I-502 FOR PTS WITH ST-SEGMENT ELEVATION ACUTE MYOCARDIAL INFARCTION, PLAVIX TO REDUCE RATE OF DEATH FROM ANY CAUSE AND THE RATE OF A COMBINED ENDPOINT OF DEATH, REINFARCTION OR STROKE. NOT KNOWN TO PERTAIN TO PTS WHO RECEIVE PRIMARY ANGIOPLASTY

I-503 TREATMENT OF MAJOR DEPRESSIVE EPISODES ASSOCIATED WITH BIPOLAR DISORDER

I-504 TREATMENT OF PATHOLOGICAL HYPERSECRETORY CONDITIONS INCLUDING ZOLLINGER-ELLISON SYNDROME

I-505 TREATMENT OF STAPHYLOCOCCUS AUREUS BLOODSTREAM INFECTIONS (BACTEREMIA), INCLUDING THOSE WITH RIGHT SIDED INFECTIVE ENDOCARDITIS, CAUSED BY METHICILLIN-SUSCEPTIBLE AND METHICILLIN-RESISTANT ISOLATES

I-506 ADJUNCTIVE THERAPY OF MYOCLONIC SEIZURES IN ADULTS AND ADOLESCENTS AGE 12 AND OVER WITH JUVENILE MYOCLONIC EPILEPSY

I-507 ADJUNCT TO DIET TO REDUCE TOTAL-C, LDL-C AND APO B LEVELS IN ADOLESCENT BOYS AND GIRLS WHO ARE AT LEAST ONE YEAR POST-MENARCHE, 10-16 YEARS OF AGE, WITH HETEROZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA

I-508 PREMENSTRUAL DYSPHONIC DISORDER

I-509 TREATMENT OF IRRITABILITY ASSOCIATED WITH AUTISTIC DISORDER

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

I-510 ADULT DERMATOFIBROSARCOMA PROTUBERANS (DFSP)

I-511 ADULT MYELODYSPLASTIC SYNDROME/MYELOPROLIFERATIVE DISEASES (MDS/MDP)

I-512 ADULT PH+ ACUTE LYMPHOBLASTIC LEUKEMIA (ALL) MONOTHERAPY

I-513 ADULT AGGRESSIVE SYSTEMIC MASTOCYTOSIS (ASM)

I-514 ADULT HYPEREOSINOPHILIC SYNDROME/CHRONIC EOSINOPHILIC LEUKEMIA (HES/CEL)

I-515 PROPHYLAXIS OF SURGICAL SITE INFECTION FOLLOWING ELECTIVE COLORECTAL SURGERY

I-516 PRIMARY GENERALIZED TONIC CLONIC SEIZURES IN ADULTS AND PEDIATRIC PATIENTS 2 YEARS OF AGE AND OLDER

I-517 TREATMENT OF MODERATE TO SEVERE PRIMARY RESTLESS LEG SYNDROME (RLS)

I-518 TREATMENT OF SHORT STATURE OR GROWTH FAILURE IN CHILDREN WITH SHOX (SHORT STATURE HOMEBOX CONTAINING GENE) DEFICIENCY WHOSE EPIPHYSES ARE NOT CLOSED

I-519 USE OF TAXOTERE (DOCETAXEL) INJECTION CONCENTRATE IN COMBINATION WITH CISPLATIN AND FLUOROURACIL FOR THE INDUCTION OF PATIENTS WITH INOPERABLE LOCALLY ADVANCED SQUAMOUS CELL CARCINOMA OF THE HEAD AND NECK (SCCHN)

I-520 USE OF EXENATIDE IN PATIENTS WITH TYPE 2 DIABETES MELLITUS WHO ARE USING A THIAZOLIDINEDIONE ALONE OR IN COMBINATION WITH METFORMIN BUT HAVE NOT ACHIEVED ADEQUATE GLYCEMIC CONTROL

I-521 TREATMENT OF PATIENTS WITH MANTLE CELL LYMPHOMA WHO HAVE RECEIVED AT LEAST 1 YEAR PRIOR THERAPY

I-522 TREATMENT OF MODERATE ACNE VULGARIS IN WOMEN AT LEAST 14 YRS OF AGE, WHO HAVE NO KNOWN CONTRAINDICATIONS TO ORAL CONTRACEPTIVE THERAPY, AND HAVE ACHIEVED MENARCHE, IF THE PATIENT DESIRES AN ORAL CONTRACEPTIVE FOR BIRTH CONTROL.

I-523 USE IN ADULT PATIENTS WITH CLINICALLY EVIDENT CORONARY HEART DISEASE TO REDUCE THE RISK OF NONFATAL MYOCARDIAL INFARCTION, FATAL AND NONFATAL STROKE, ANGINA, REVASCULARIZATION PROCEDURES AND HOSPITALIZATION FOR CONGESTIVE HEART FAILURE

I-524 GENERALIZED ANXIETY DISORDER (GAD)

I-525 USE OF 0.5MG/0.1MG FOR PREVENTION OF POST-MENOPAUSAL OSTEOPOROSIS

I-526 TREATMENT OF HYPONATREMIA IN HOSPITALIZED PATIENTS

I-527 ADJUNCTIVE THERAPY IN THE TREATMENT OF PRIMARY GENERALIZED TONIC-CLONIC SEIZURES IN ADULTS AND CHILDREN 6 YEARS OF AGE AND OLDER WITH IDIOPATHIC GENERALIZED EPILEPSY

I-528 TREATMENT OF MODERATE TO SEVERE VAGINAL DRYNESS AND PAIN WITH INTERCOURSE, SYMPTOMS OF VULVAR AND VAGINAL ATROPHY ASSOCIATED WITH MENOPAUSE

I-529 TREATMENT OF DEMENTIA OF THE ALZHEIMER'S TYPE IN PATIENTS WITH SEVERE ALZHEIMER'S DISEASE

I-530 PREVENTION OF EXERCISE-INDUCED BRONCHOCONSTRICTION IN PATIENTS 15 YEARS OF AGE AND OLDER

I-531 MAINTENANCE TREATMENT OF SCHIZOPHRENIA

I-532 TREATMENT OF BACTERIAL VAGINOSIS IN NON-PREGNANT FEMALES

I-533 ACUTE ST-SEGMENT ELEVATION MYOCARDIAL INFARCTION (STEMI)

I-534 EXTENDED TREATMENT OF SYMPTOMATIC VENOUS THROMBOEMBOLISM (VTE) AND/OR PULMONARY EMBOLISM TO REDUCE THE RECURRENCE OF VTE IN PATIENTS WITH CANCER

I-535 MANAGEMENT OF FIBROMYALGIA

I-536 FOR THE TREATMENT OF SHORT STATURE IN CHILDREN WITH NOONAN SYNDROME

I-537 LONG TERM TREATMENT OF PANIC DISORDER

I-538 SHORT TERM TREATMENT OF PANIC DISORDER

I-539 REDUCTION IN RISK OF INVASIVE BREAST CANCER IN POSTMENOPAUSAL WOMEN WITH OSTEOPOROSIS OR AT HIGH RISK FOR INVASIVE BREAST CANCER

I-540 TREATMENT OF SCHIZOPHRENIA IN ADOLESCENTS AGES 13-17

I-541 TREATMENT OF BIPOLAR I DISORDER IN CHILDREN AGES 10-12 AND ADOLESCENTS AGES 13-17

I-542 EXPANSION OF PATIENT POPULATION FOR HEAD AND NECK CANCER FROM "INOPERABLE" PATIENTS TO ALL PATIENTS

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

- I-543 USE IN COMBINATION WITH CISPLATIN AND FLUOROURACIL FOR THE INDUCTION TREATMENT OF PATIENTS WITH LOCALLY ADVANCED SQUAMOUS CELL CARCINOMA OF THE HEAD AND NECK (SCCHN)
- I-544 ADJUNCTIVE THERAPY OF MYOCLONIC SEIZURES IN ADULTS AND ADOLESCENTS AGE 16 AND OVER WITH JUVENILE MYOCLONIC EPILEPSY
- I-545 ADJUNCTIVE TREATMENT TO TREAT PATIENTS WITH MAJOR DEPRESSIVE DISORDER
- I-546 TREATMENT OF UNRESECTABLE HEPATOCELLULAR CARCINOMA
- I-547 ADJUNCTIVE THERAPY TO DIET TO SLOW THE PROGRESSION OF ARTEROSCLEROSIS IN ADULT PATIENTS AS PART OF A TREATMENT STRATEGY TO LOWER TOTAL-C AND LDL-C TO TARGET LEVELS
- I-548 SEASONAL ALLERGIC RHINITIS IN PATIENTS 6 THROUGH LESS THAN 12 YEARS OF AGE
- I-549 USE OF AVALIDE TABLETS AS INITIAL THERAPY IN PATIENTS WHO ARE LIKELY TO NEED MULTIPLE DRUGS TO ACHIEVE THEIR BLOOD PRESSURE GOALS
- I-550 TREATMENT OF HYPERTENSION IN PEDIATRIC PATIENTS 6-16 YEARS OF AGE
- I-551 TREATMENT OF SHORT STATURE IN CHILDREN WITH TURNER'S SYNDROME
- I-552 ADJUNCTIVE TREATMENT FOR RADIOIODINE ABLATION OF THYROID TISSUE REMNANTS IN PATIENTS WHO HAVE UNDERGONE THYROIDECTOMY FOR WELL-DIFFERENTIATED THYROID CANCER AND WHO DO NOT HAVE EVIDENCE OF METASTATIC THYROID CANCER
- I-553 FOR USE AS AN ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS
- I-554 TREATMENT OF PATIENTS WITH CANDIDEMIA, ACUTE DISSEMINATED CANDIDIASIS, CANDIDA PERITONITIS AND ABSCESES
- I-555 TREATMENT OF ACUTE MANIC OR MIXED EPISODES ASSOCIATED WITH BIPOLAR I DISORDER IN PEDIATRIC PATIENTS AGED 10-17 YEARS
- I-556 PREVENTION OF POST OPERATIVE NAUSEA AND VOMITING FOR UP TO 24 HOURS FOLLOWING SURGERY
- I-557 USE OF AMITIZA (LUBIPROSTONE) 8 MCG TWICE DAILY FOR TREATMENT OF IRRITABLE BOWEL SYNDROME WITH CONSTIPATION IN WOMEN GREATER THAN OR EQUAL TO 18 YEARS OLD
- I-558 MAINTENANCE TREATMENT OF AIRFLOW OBSTRUCTION AND REDUCING EXACERBATIONS IN PATIENTS WITH CHRONIC OBSTRUCTIVE PULMONARY DISEASE (COPD) INCLUDING CHRONIC BRONCHITIS AND EMPHYSEMA
- I-559 ADJUNCTIVE THERAPY ADDED TO LITHIUM OR VALPROATE IN SHORT TERM TREATMENT OF BIPOLAR DISORDER, MANIC OR MIXED
- I-560 MAINTENANCE TREATMENT FOR BIPOLAR I DISORDER, AS ADJUNCTIVE THERAPY TO LITHIUM OR DIVALPROEX
- I-561 LONG-TERM TREATMENT OF SOCIAL ANXIETY DISORDER
- I-562 MAINTENANCE TREATMENT OF ATTENTION-DEFICIT HYPERACTIVITY DISORDER (ADHD) IN CHILDREN AND ADOLESCENTS
- I-563 ADJUNCTIVE THERAPY IN THE TREATMENT OF PRIMARY GENERALIZED TONIC-CLONIC SEIZURES IN ADULTS AND CHILDREN 16 YEARS OF AGE AND OLDER WITH IDIOPATHIC GENERALIZED EPILEPSY
- I-564 TREATMENT OF PATIENTS WITH MULTIPLE MYELOMA
- I-565 USE OF DUTASTERIDE IN COMBINATION WITH TAMSULOSIN FOR THE TREATMENT OF SYMPTOMATIC BENIGN PROSTATIC HYPERPLASIA (BPH)
- I-566 MANAGEMENT OF FIBROMYALGIA
- I-567 INITIAL THERAPY IN PATIENTS LIKELY TO NEED MULTIPLE DRUGS TO ACHIEVE THEIR BLOOD PRESSURE GOALS
- I-568 USE OF APTIVUS, CO-ADMINISTERED W/RITONAVIR, FOR COMBINATION ANTIRETROVIRAL TREATMENT OF HIV-1 INFECTED PED (AGE 2-18 YRS) PATIENTS WHO ARE TREATMENT-EXPERIENCED AND INFECTED W/HIV-1 STRAINS RESISTANT TO MORE THAN ONE PROTEASE INHIBITOR
- I-569 TREATMENT OF CHRONIC HEPATITIS B
- I-570 TREATMENT OF CHICKEN POX IN IMMUNOCOMPETENT PEDIATRIC PATIENTS 2 TO <18 YEARS OF AGE
- I-571 NON-SMALL CELL LUNG CANCER IN COMBINATION WITH CISPLATIN AND AS SINGLE AGENT FOR NON-SQUAMOUS NON-SMALL CELL LUNG CANCER

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

I-572	TREATMENT OF GROWTH FAILURE IN CHILDREN BORN SMALL FOR GESTATIONAL AGE (SGA) WITH NO CATCH-UP BY AGE 2-4 YRS.
I-573	TO TREAT PATIENTS WITH PRIMARY DYSBETALIPOPROTEINEMIA (FREDRICKSON TYPE III HYPERLIPOPROTEINEMIA) AS AN ADJUNCT TO DIET
I-574	MONOTHERAPY IN THE TREATMENT OF BIPOLAR DEPRESSION
I-575	MONOTHERAPY IN THE TREATMENT OF BIPOLAR MANIA
I-576	ADJUNCTIVE THERAPY IN THE TREATMENT OF BIPOLAR MANIA
I-577	SEDATION OF NON-INTUBATED PATIENTS PRIOR TO AND/OR DURING SURGICAL AND OTHER PROCEDURES
I-578	EXPANSION OF INDICATION TO INCLUDE TREATMENT OF HIV IN TREATMENT NAIVE ADULTS
I-579	TREATMENT OF MODERATE TO SEVERE DYSpareunia, A SYMPTOM OF VULVAR AND VAGINAL ATROPHY, DUE TO MENOPAUSE AND NEW TWICE WEEKLY DOSING REGIMEN FOR THIS INDICATION
I-580	INDOLENT B-CELL NON-HODGKINS LYMPHOMA (NHL) THAT HAS PROGRESSED DURING OR WITHIN SIX MONTHS OF TREATMENT WITH RITUXIMAB OR A RITUXIMAB CONTAINING REGIMEN
I-581	TREATMENT TO INCREASE BONE MASS IN MEN WITH OSTEOPOROSIS
I-582	TREATMENT OF CHRONIC OBSTRUCTIVE PULMONARY DISEASE
I-583	ADJUVANT TREATMENT OF ADULT PATIENTS FOLLOWING COMPLETE GROSS RESECTION OF KIT (CD117) POSITIVE GASTROINTESTINAL STROMAL TUMORS (GIST)
I-584	TREATMENT AND PREVENTION OF GLUCOCORTICOID-INDUCED OSTEOPOROSIS IN PATIENTS EXPECTED TO BE ON GLUCOCORTICIDS FOR AT LEAST 12 MONTHS
I-585	TREATMENT OF SHORT STATURE IN PEDIATRIC PATIENTS SMALL FOR GESTATIONAL AGE WHO DO NOT MANIFEST CATCH UP GROWTH BY AGE 2 TO 4 YEARS
I-586	COMMUNITY ACQUIRED BACTERIAL PNEUMONIA
I-587	ADDITIONAL PATHOGENS TO COMPLICATED SKIN AND SKIN STRUCTURE INFECTIONS INDICATION
I-588	ADDITIONAL PATHOGENS TO COMPLICATED INTRA-ABDOMINAL INFECTIONS INDICATION
I-589	TREATMENT OF TREATMENT RESISTANT DEPRESSION (TRD) IN COMBINATION WITH OLANZAPINE
I-590	ACUTE TREATMENT OF DEPRESSIVE EPISODES ASSOCIATED WITH BIPOLAR DISORDER (IN COMBINATION WITH OLANZAPINE)
I-591	TREATMENT OF TREATMENT RESISTANT DEPRESSION (TRD) IN COMBINATION WITH FLUOXETINE
I-592	ACUTE TREATMENT OF DEPRESSIVE EPISODES ASSOCIATED WITH BIPOLAR DISORDER (IN COMBINATION WITH FLUOXETINE)
I-593	TREATMENT OF TREATMENT RESISTANT DEPRESSION (TRD)
I-594	INDICATION EXPANDED TO INCLUDE PATIENTS WHO HAVE EXPERIENCED A FIRST CLINICAL EPISODE AND HAVE MRI FEATURES CONSISTENT WITH MULTIPLE SCLEROSIS
I-595	PREVENTION OF OSTEOPOROSIS IN POSTMENOPAUSAL WOMEN
I-596	USE AS ADJUNCTIVE THERAPY WITH LITHIUM OR VALPROATE FOR THE MAINTENANCE TREATMENT OF BIPOLAR I DISORDER
I-597	MONOTHERAPY FOR THE MAINTENANCE TREATMENT OF BIPOLAR I DISORDER
I-598	TREATMENT OF PULMONARY ARTERIAL HYPERTENSION INDICATION EXPANDED TO INCLUDE DELAY IN CLINICAL WORSENING
I-599	PREVENTION AND TREATMENT OF SECONDARY HYPERPARATHYROIDISM ASSOCIATED WITH CHRONIC KIDNEY DISEASE (CKD) STAGE 5 IN PATIENTS ON HEMODIALYSIS OR PERITONEAL DIALYSIS
I-600	FOR USE AS INITIAL THERAPY IN PATIENTS WHO ARE LIKELY TO NEED MULTIPLE DRUGS TO ACHIEVE THEIR BLOOD PRESSURE GOALS
I-601	MAINTENANCE TREATMENT IN PATIENTS WITH ADVANCED OR METASTATIC NONSMALL CELL LUNG CANCER WHOSE DISEASE HAS NOT PROGRESSED AFTER FOUR CYCLES OF PLATINUM-BASED FIRST LINE CHEMOTHERAPY
I-602	TREATMENT OF MEN AND WOMEN WITH OSTEOPOROSIS ASSOCIATED WITH SUSTAINED SYSTEMIC GLUCOCORTICOID THERAPY AT HIGH RISK FOR FRACTURE
I-603	GOUT FLARES
I-604	PREVENTION OF CMV DISEASE IN KIDNEY AND HEART TRANSPLANT PATIENTS 4 MONTHS TO 16

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

YEARS AT HIGH RISK

- I-605 ADJUNCT TO MOOD STABILIZERS AND/OR ANTIDEPRESSANTS FOR SCHIZOAFFECTIVE DISORDER
- I-606 TREATMENT OF SCHIZOAFFECTIVE DISORDER AS MONOTHERAPY
- I-607 INDICATION EXPANDED TO INCLUDE TREATMENT OF PULMONARY ARTERIAL HYPERTENSION (WHO GROUP I) IN PATIENTS WITH CLASS II SYMPTOMS
- I-608 REDUCE LDL-C LEVELS IN BOYS AND POSTMENARCHAL GIRLS, 10 TO 17 YEARS OF AGE, WITH HETEROZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA AS MONOTHERAPY OR IN COMBINATION WITH A STATIN AFTER FAILING AN ADEQUATE TRIAL OF DIET THERAPY
- I-610 TREATMENT OF HEAVY MENSTRUAL BLEEDING FOR WOMEN WHO CHOOSE TO USE INTRAUTERINE CONTRACEPTION AS THEIR METHOD OF CONTRACEPTION
- I-611 TREATMENT OF HETEROZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA IN ADOLESCENT BOYS AND POSTMENARCHAL GIRLS, AGES 10 TO 17 YEARS, WITH A RECOMMENDATION DOSING RANGE OF 5 TO 20 MG ONCE DAILY
- I-612 MICARDIS 80 MG FOR REDUCTION OF THE RISK OF MYOCARDIAL INFARCTION, STROKE, OR DEATH FROM CARDIOVASCULAR CAUSES IN PATIENTS 55 YEARS OF AGE OR OLDER AT HIGH RISK OF DEVELOPING MAJOR CARDIOVASCULAR EVENTS WHO ARE UNABLE TO TAKE ACE INHIBITORS
- I-613 MILD TO MODERATE ATOPIC DERMATITIS IN PATIENTS 3 MONTHS OF AGE TO LESS THAN 18 YEARS OF AGE
- I-614 SHORT TERM TREATMENT OF EROSIIVE ESOPHAGITIS ASSOCIATED WITH GERD IN PEDIATRIC PATIENTS AGES FIVE YEARS AND OLDER
- I-615 MAINTENANCE TREATMENT OF BIPOLAR DISORDER AS AN ADJUNCT TO LITHIUM OR VALPROATE
- I-616 TREATMENT OF IRRITABILITY ASSOCIATED WITH AUTISTIC DISORDER IN PEDIATRIC PATIENTS AGES 6-17 YEARS OF AGE
- I-617 MAINTENANCE OF GENERALIZED ANXIETY DISORDER (GAD)
- I-618 ADJUNCTIVE THERAPY IN THE TREATMENT OF MAJOR DEPRESSIVE DISORDER (MDD)
- I-619 INTRAVENOUS CONTRAST ENHANCED COMPUTER TOMOGRAPHY OF THE HEAD AND BODY
- I-620 FOR USE IN COMBINATION WITH LETROZOLE FOR THE TREATMENT OF POSTMENOPAUSAL WOMEN WITH HORMONE RECEPTOR POSITIVE METASTATIC BREAST CANCER THAT OVEREXPRESSES THE HER2 RECEPTOR FOR WHOM HORMONAL THERAPY IS INDICATED
- I-621 PRIMARY PREVENTION OF CARDIOVASCULAR DISEASE, BASED ON THE RESULTS OF JUSTIFICATION FOR THE USE OF STATINS IN PRIMARY PREVENTION; AN INTERVENTION TRIAL EVALUATING ROSUVASTATIN (JUPITER)
- I-622 ADJUNCTIVE THERAPY FOR PRIMARY GENERALIZED TONIC-CLONIC SEIZURES IN PATIENTS THIRTEEN YEARS OF AGE AND OLDER
- I-623 TREATMENT OF SIGNS AND SYMPTOMS OF ADVANCED IDIOPATHIC PARKINSON'S DISEASE
- I-624 MAINTENANCE TREATMENT OF PATIENTS WITH LOCALLY ADVANCED OR METASTATIC NON-SMALL CELL LUNG CANCER WHOSE DISEASE HAS NOT PROGRESSED AFTER FOUR CYCLES OF PLATINUM-BASED FIRST-LINE CHEMOTHERAPY
- I-625 PANCREATIC INSUFFICIENCY DUE TO CHRONIC PANCREATITIS AND PANCREATECTOMY
- I-626 RELIEF OF NASAL CONGESTION ASSOCIATED WITH SEASONAL ALLERGIC RHINITIS IN ADULTS AND PEDIATRIC PATIENTS 2 YEARS OF AGE AND OLDER
- I-627 TREATMENT OF NEWLY DIAGNOSED ADULT PATIENTS WITH PHILADELPHIA CHROMOSOME POSITIVE CHRONIC MYELOID LEUKEMIA (PH & CML) IN CHRONIC PHASE.
- I-628 MAINTENANCE TREATMENT OF SCHIZOPHRENIA IN ADULTS
- I-629 ADJUNCTIVE THERAPY WITH EITHER LITHIUM OR VALPROATE FOR THE ACUTE TREATMENT OF MANIC OR MIXED EPISODES ASSOCIATED WITH BIPOLAR I DISORDER
- I-630 TREATMENT OF PATIENTS WITH SUBEPENDYMAL GIANT CELL ASTROCYTOMA (SEGA) ASSOCIATED WITH TUBEROUS SCLEROSIS (TS) WHO REQUIRE THERAPEUTIC INTERVENTION BUT ARE NOT CANDIDATES FOR CURATIVE SURGICAL RESECTION.
- I-631 PREVENTION OF RELAPSE TO OPIOID DEPENDENCE FOLLOWING OPIOID DETOXIFICATION
- I-632 MANAGEMENT OF CHRONIC MUSCULOSKELETAL PAIN
- I-633 MAINTENANCE TREATMENT OF BIPOLAR I DISORDER AS AN ADJUNCT TO LITHIUM OR VALPROATE
- I-634 TREATMENT OF SEVERE HYPERCALCEMIA IN PATIENTS WITH PRIMARY HYPERPARATHYROIDISM WHO ARE UNABLE TO UNDERGO PARATHYROIDECTOMY

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

I-635	ADJUNCTIVE TREATMENT WITH LONG-ACTING ORAL PSYCHOSTIMULANTS FOR THE TREATMENT OF ATTENTION DEFICIT HYPERACTIVITY DISORDER (ADHD)
I-636	TREATMENT OF EXTERNAL GENITAL AND PERIANAL WARTS/CONDYLOMA ACUMINATA IN PATIENTS 12 YEARS OR OLDER
I-637	USE IN COMBINATION CHEMOTHERAPY WITH 5-FLUOROURACIL IN THE PALLIATIVE TREATMENT OF PATIENTS WITH ADVANCED METASTATIC COLORECTAL CANCER
I-638	FOR PATIENTS WITH PROGRESSIVE NEUROENDOCRINE TUMORS OF PANCREATIC ORIGIN (PNET) THAT ARE UNRESECTABLE, LOCALLY ADVANCED, OR METASTATIC.
I-639	TREATMENT OF PROGRESSIVE, WELL-DIFFERENTIATED PANCREATIC NEUROENDOCRINE TUMORS IN PATIENTS WITH UNRESECTABLE, LOCALLY ADVANCED, OR METASTATIC DISEASE
I-640	MAINTENANCE OF REMISSION OF ULCERATIVE COLITIS
I-641	TREATMENT OF THE SIGNS AND SYMPTOMS OF BENIGN PROSTATIC HYPERPLASIA (BPH)
I-642	TREATMENT OF ERECTILE DYSFUNCTION (ED) AND THE SIGNS AND SYMPTOMS OF BENIGN PROSTATIC HYPERPLASIA (BPH)
I-643	REDUCE THE RISK OF STROKE AND SYSTEMIC EMBOLISM IN PATIENTS WITH NONVALVULAR ATRIAL FIBRILLATION.
I-644	MONOTHERAPY IN PATIENTS 13 YEARS OF AGE AND OLDER WITH PARTIAL SEIZURES WHO ARE RECEIVING THERAPY WITH A SINGLE ANTIEPILEPTIC DRUG (AED)
I-645	MAINTENANCE TREATMENT OF ATTENTION DEFICIT HYPERACTIVITY DISORDER (ADHD) IN ADULTS
I-646	SIGNS AND SYMPTOMS OF ADVANCED PARKINSON'S DISEASE (APD)
I-647	SIGNS AND SYMPTOMS OF MODERATE TO SEVERE PRIMARY RESTLESS LEGS SYNDROME (RLS)
I-648	TREATMENT OF HEAVY MENSTRUAL BLEEDING IN WOMEN WITHOUT ORGANIC PATHOLOGY WHO CHOOSE TO USE AN ORAL CONTRACEPTIVE AS THEIR METHOD OF CONTRACEPTION
I-649	TREATMENT OF PATIENTS WITH ADVANCED SOFT TISSUE SARCOMA (STS) WHO HAVE RECEIVED PRIOR CHEMOTHERAPY
I-650	TREATMENT OF ADULTS WITH RENAL ANGIOMYOLIPOMA AND TUBEROUS SCLEROSIS COMPLEX (TSC), NOT REQUIRING IMMEDIATE SURGERY
I-651	MANAGEMENT OF NEUROPATHIC PAIN ASSOCIATED WITH SPINAL CORD INJURY
I-652	MANAGEMENT OF POSTHERPETIC NEURALGIA
I-653	TREATMENT OF ENDOGENOUS ANTERIOR UVEITIS
I-654	MAGNETIC RESONANCE ANGIOGRAPHY (MRA) TO EVALUATE ADULTS WITH KNOWN OR SUSPECTED RENAL OR AORTO-ILIO-FEMORAL OCCLUSIVE VASCULAR DISEASE
I-655	TREATMENT OF POSTMENOPAUSAL WOMEN WITH ADVANCED HORMONE RECEPTOR-POSITIVE, HER2-NEGATIVE BREAST CANCER (ADVANCED HR+BC) IN COMBINATION WITH EXEMESTANE, AFTER FAILURE OF TREATMENT WITH LETROZOLE OR ANASTROZOLE
I-656	MANAGEMENT OF NEUROPATHIC PAIN ASSOCIATED WITH DIABETIC PERIPHERAL NEUROPATHY (DPN) IN ADULTS WHEN A CONTINUOUS, AROUND-THE-CLOCK OPIOID ANALGESIC IS NEEDED FOR AN EXTENDED PERIOD OF TIME
I-657	PLAQUE PSORIASIS OF THE SCALP
I-658	FIRST-LINE TREATMENT OF LOCALLY ADVANCED OR METASTATIC NON-SMALL CELL LUNG CANCER, IN COMBINATION WITH CARBOPLATIN, IN PATIENTS WHO ARE NOT CANDIDATES FOR CURATIVE SURGERY OR RADIATION THERAPY
I-659	PLAQUE PSORIASIS OF THE BODY
I-660	TREATMENT OF DEEP VEIN THROMBOSIS
I-661	TREATMENT OF PULMONARY EMBOLISM
I-662	REDUCTION IN RISK FOR DEEP VEIN THROMBOSIS AND THE REDUCTION IN RISK FOR PULMONARY EMBOLISM
I-663	IN COMBINATION WITH PREDNISONE FOR THE TREATMENT OF PATIENTS WITH METASTATIC CASTRATION-RESISTANT PROSTATE CANCER
I-664	TREATMENT OF THROMBOCYTOPENIA IN PATIENTS WITH CHRONIC HEPATITIS C TO ALLOW THE INITIATION AND MAINTENANCE OF INTERFERON-BASED THERAPY
I-665	TREATMENT OF CHRONIC IRON OVERLOAD IN PATIENTS 10 YRS OF AGE AND OLDER WITH (NTDT)SYNDROMES AND WITH A (LIC) OF AT LEAST 5 MG OF IRON PER GRAM OF LIVER DRY WEIGHT (MG FE/G DW) AND SERUM FERRITIN GREATER THAN 300MCG/L

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

I-666	TREATMENT OF PEDIATRIC PATIENTS WITH NEWLY DIAGNOSED PHILADELPHIA CHROMOSOME-POSITIVE ACUTE LYMPHOBLASTIC LEUKEMIA (PH+ALL) IN COMBINATION WITH CHEMOTHERAPY
I-667	TREATMENT OF PATIENTS WITH LOCALLY ADVANCED, UNRESECTABLE OR METASTATIC GASTROINTESTINAL STROMAL TUMOR (GIST) WHO HAVE BEEN PREVIOUSLY TREATED WITH IMATINIB MESYLATE AND SUNITINIB MALATE
I-668	PROPHYLAXIS OF ALLOGRAFT REJECTION IN ADULT PATIENTS RECEIVING A LIVER TRANSPLANT
I-669	SCINTIGRAPHIC ASSESSMENT OF SYMPATHETIC INNERVATION OF THE MYOCARDIUM BY MEASUREMENT OF THE HEART TO MEDIASTINUM (H/M) RATIO OF RADIOACTIVITY UPTAKE IN PATIENTS WITH NYHA CLASS II OR CLASS III HEART FAILURE AND LVEF LESS THAN 35%
I-670	TREATMENT OF OPIOID-INDUCED CONSTIPATION (OIC) IN ADULTS WITH CHRONIC, NON-CANCER PAIN
I-671	FIRSTLINE TREATMENT OF PATIENTS WITH METASTATIC NON- SMALL CELL LUNG CANCER (NSCLC) WHOSE TUMORS HAVE EPIDERMAL GROWTH FACTOR RECEPTOR (EGFR) EXON 19 DELETIONS OR EXON 21 (L858R) SUBSTITUTION MUTATIONS AS DETECTED BY AN FDA-APPROVED TEST
I-672	USE IN PATIENTS WITH MANTLE CELL LYMPHOMA WHOSE DISEASE HAS RELAPSED OR PROGRESSED AFTER TWO PRIOR THERAPIES, ONE OF WHICH INCLUDED BORTEZOMIB
I-673	TREATMENT OF HOSPITAL-ACQUIRED BACTERIAL PNEUMONIA/VENTILATOR-ASSOCIATED BACTERIAL PNEUMONIA (HABP/VABP) CAUSED BY SUSCEPTIBLE ISOLATES OF S. AUREUS (INCLUDING METHICILLIN-SUSCEPTIBLE AND RESISTANT ISOLATES) WHEN ALTERNATIVE TREATMENTS ARE NOT SUITABLE
I-674	TREATMENT OF PATIENTS WITH DEPRESSIVE EPISODES ASSOCIATED WITH BIPOLAR I DISORDER (BIPOLAR DEPRESSION) AS MONOTHERAPY AND AS ADJUNCTIVE THERAPY WITH LITHIUM OR VALPROATE
I-675	MAINTENANCE TREATMENT OF MAJOR DEPRESSIVE DISORDER
I-676	FIRST-LINE TREATMENT OF PATIENTS WITH METASTATIC ADENOCARCINOMA OF THE PANCREAS, IN COMBINATION WITH GEMCITABINE
I-677	TREATMENT OF PATIENTS WITH LOCALLY RECURRENT OR METASTATIC, PROGRESSIVE, DIFFERENTIATED THYROID CARCINOMA (DTC) THAT IS REFRACTORY TO RADIOACTIVE IODINE TREATMENT
I-678	TRAMETINIB, IN COMBINATION WITH DABRAFENIB, FOR THE TREATMENT OF PATIENTS WITH UNRESECTABLE OR METASTATIC MELANOMA WITH BRAF V600E OR V600K MUTATIONS AS DETECTED BY AN FDA-APPROVED TEST
I-679	RISK REDUCTION OF REBLEEDING OF GASTRIC OR DUODENAL ULCERS FOLLOWING THERAPEUTIC ENDOSCOPY IN ADULTS
I-680	TREATMENT OF PATIENTS WITH CHRONIC LYMPHOCYTIC LEUKEMIA (CLL) WHO HAVE RECEIVED AT LEAST ONE PRIOR THERAPY
I-681	PROPHYLAXIS OF DEEP VEIN THROMBOSIS (DVT) WHICH MAY LEAD TO PULMONARY EMBOLISM (PE), IN ADULT PATIENTS WHO HAVE UNDERGONE HIP OR KNEE REPLACEMENT
I-682	TREATMENT OF DEEP VEIN THROMBOSIS (DVT) AND PULMONARY EMBOLISM (PE) IN PATIENTS WHO HAVE BEEN TREATED WITH A PARENTERAL ANTICOAGULANT FOR 5-10 DAYS
I-683	TO REDUCE THE RISK OF RECURRENCE OF DVT AND PE IN PATIENTS WHO HAVE BEEN PREVIOUSLY TREATED
I-684	PREVENTION OF ACUTE NAUSEA AND VOMITING ASSOCIATED WITH INITIAL AND REPEAT COURSES OF EMETOGENIC CANCER CHEMOTHERAPY, INCLUDING HIGHLY EMETOGENIC CANCER CHEMOTHERAPY IN PEDIATRIC PATIENTS AGED 1 MONTH TO LESS THAN 17 YEARS
I-685	EXPANDED INDICATION OF RASAGILINE AS AN ADD-ON THERAPY TO STABLE DOSES OF DOPAMINE AGONISTS IN THE TREATMENT OF EARLY PARKINSON'S DISEASE
I-686	INDICATED FOR THE TREATMENT OF DIABETIC MACULAR EDEMA IN PATIENTS WHO ARE PSEUDOPHAKIC OR ARE PHAKIC AND SCHEDULED FOR CATARACT SURGERY
I-687	GUIDING SENTINEL LYMPH NODE BIOPSY, USING A HAND-HELD GAMMA COUNTER IN PATIENTS WITH CLINICALLY NODE NEGATIVE SQUAMOUS CELL CARCINOMA OF THE ORAL CAVITY
I-688	GADAVIST IS INDICATED WITH MRI TO DETECT THE PRESENCE AND EXTENT OF MALIGNANT BREAST DISEASE
I-689	TREATMENT OF PATIENTS WITH CHRONIC LYMPHOCYTIC LEUKEMIA (CLL) WITH 17P DELETION
I-690	INDICATED FOR THE TREATMENT OF DEEP VEIN THROMBOSIS (DVT)
I-691	INDICATED TO REDUCE THE RISK OF RECURRENT DEEP VEIN THROMBOSIS (DVT) AND

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

PULMONARY EMBOLISM (PE) FOLLOWING INITIAL THERAPY

I-692 INDICATED FOR MANAGEMENT OF OSTEOARTHRITIS PAIN.

I-693 TREATMENT OF PATIENTS WITH METASTATIC CASTRATION-RESISTANT PROSTATE CANCER (CRPC)

I-694 TREATMENT OF PATIENTS WITH MODERATE TO SEVERE PLAQUE PSORIASIS WHO ARE CANDIDATES FOR PHOTOTHERAPY OR SYSTEMIC THERAPY

I-695 REVISED INDICATION FOR BORTEZOMIB IN THE TREATMENT OF PATIENTS WITH MANTLE CELL LYMPHOMA

I-696 USE AS MONOTHERAPY IN THE TREATMENT OF PARTIAL-ONSET SEIZURES IN PATIENTS WITH EPILEPSY AGE 17 YEARS AND OLDER

I-697 FOR USE IN COMBINATION WITH SOFOSBUVIR FOR THE TREATMENT OF PATIENTS WITH CHRONIC HEPATITIS C VIRUS GENOTYPE 1 INFECTION

I-698 SCHIZOAFFECTIVE DISORDER AS MONOTHERAPY AND AS AN ADJUNCT TO MOOD STABILIZERS OR ANTIDEPRESSANTS

I-699 FOR TREATMENT OF PATIENTS WITH POLYCYTHEMIA VERA WHO HAVE HAD AN INADEQUATE RESPONSE TO OR ARE INTOLERANT OF HYDROXYUREA

I-700 TREATMENT OF PEDIATRIC PATIENTS WITH TOURETTE'S DISORDER (6-18 YEARS)

I-701 FOR THE TREATMENT OF PATIENTS WITH UNRESECTABLE, WELL-OR MODERATELY-DIFFERENTIATED, LOCALLY ADVANCED OR METASTATIC GASTROENTEROPANCREATIC NEUROENDOCRINE TUMORS (GEP-NETS) TO IMPROVE PROGRESSION FREE SURVIVAL

I-702 FOR THE TREATMENT OF PATIENTS WITH WALDENSTROM MACROGLOBULINEMIA

I-703 MODERATE TO SEVERE BINGE EATING DISORDER (BED)

I-704 EXPANDED INDICATION TO INCLUDE PATIENTS WHO ARE VIROLOGICALLY-SUPPRESSED (HIV-1 RNA <50 COPIES/ML) ON A STABLE ANTIRETROVIRAL REGIMEN FOR AT LEAST 6 MONTHS WITH NO HISTORY OF TREATMENT FAILURE IN ORDER TO REPLACE THEIR CURRENT REGIMEN

I-705 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS 6 YEARS AND OLDER WHO HAVE AN R117H MUTATION IN THE CFTR GENE

I-706 EXPANDED INDICATION FOR THE TREATMENT OF PATIENTS WITH MULTIPLE MYELOMA

I-707 POMALYST, IN COMBINATION WITH DEXAMETHASONE, IS INDICATED FOR PATIENTS WITH MULTIPLE MYELOMA WHO HAVE RECEIVED AT LEAST 2 PRIOR THERAPIES AND HAVE DEMONSTRATED DISEASE PROGRESSION ON OR WITHIN 60 DAYS OF COMPLETION OF THE LAST THERAPY

I-708 DAILY TREATMENT OF ASTHMA IN PATIENTS AGED 18 YEARS AND OLDER

I-709 TREATMENT OF IRRITABLE BOWEL SYNDROME WITH DIARRHEA (IBS-D) IN ADULTS

I-710 ADJUNCTIVE THERAPY FOR THE TREATMENT OF PRIMARY GENERALIZED TONIC-CLONIC (PG TC) SEIZURES IN PATIENTS WITH EPILEPSY 12 YEARS OF AGE OR OLDER.

I-711 INCLUSION OF PEDIATRIC PATIENTS AGES 6 YRS AND OLDER FOR THE TREATMENT OF THROMBOCYTOPENIA IN PATIENTS WITH CHRONIC ITP WHO HAVE HAD AN INSUFFICIENT RESPONSE TO CORTICOSTEROIDS, IMMUNOGLOBULINS, OR SPLENECTOMY.

I-712 EXPANDED INDICATION FOR USE IN COMBINATION WITH LENALIDOMIDE AND DEXAMETHASONE FOR THE TREATMENT OF PATIENTS WITH RELAPSED MULTIPLE MYELOMA WHO HAVE RECEIVED ONE TO THREE PRIOR LINES OF THERAPY

I-713 REVISIONS TO THE LABELING TO PERMIT THE USE OF ZUBSOLV AS INITIAL ("INDUCTION") TREATMENT OF OPIOID DEPENDENCE

I-714 EXTENDS THE 2011 APPROVAL OF BRILINTA FOR USE BEGINNING WITH ACS TO USE BEGINNING MORE REMOTE FROM MYOCARDIAL INFARCTION

I-715 FOR THE ADDITION OF THE INDICATION FOR MONOTHERAPY TREATMENT IN PARTIAL-ONSET SEIZURES IN ADULTS.

I-716 REVISED INDICATION TO INCLUDE LANGUAGE ABOUT THE BENEFITS OF USING LETAIRIS IN COMBINATION WITH TADALAFIL TO REDUCE THE RISK OF DISEASE PROGRESSION AND HOSPITALIZATION FOR WORSENING PAH AND TO IMPROVE EXERCISE ABILITY, BASED ON THE AMBITION STUDY

I-717 EXPANDED INDICATION TO INCLUDE THE TREATMENT OF CHRONIC HEPATITIS C GENOTYPE 4

I-718 EXPANDED INDICATION TO INCLUDE SUBJECTS INFECTED WITH CHRONIC HEPATITIS C, GENOTYPE 6 VIRUS INFECTION BASED UPON THE RESULTS OF THE ELECTRON- 2 STUDY

I-719 EXPANDED INDICATION TO INCLUDE THE TREATMENT OF SUBJECTS WITH GENOTYPE 5 CHRONIC HEPATITIS C VIRUS INFECTION BASED ON THE RESULTS FROM STUDY GS-US-337-119.

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

- I-720 EXPANDED INDICATION TO INCLUDE TREATMENT OF GENOTYPE 4, CHRONIC HEPATITIS C VIRUS INFECTION BASED UPON THE RESULTS FROM STUDIES ION-4 AND GS-US-337-119.
- I-721 TREATMENT OF PATIENTS WITH UNRESECTABLE OR METASTATIC LIPOSARCOMA WHO HAVE RECEIVED A PRIOR ANTHRACYCLINE-CONTAINING REGIMEN.
- I-722 REVISED INDICATION FOR USE IN COMBINATION WITH DEXAMETHASONE OR WITH LENALIDOMIDE PLUS DEXAMETHASONE FOR THE TREATMENT OF PATIENTS WITH RELAPSED OR REFRACTORY MULTIPLE MYELOMA WHO HAVE RECEIVED ONE TO THREE LINES OF THERAPY.
- I-723 AS A SINGLE AGENT FOR THE TREATMENT OF PATIENTS WITH RELAPSED OR REFRACTORY MULTIPLE MYELOMA WHO HAVE RECEIVED ONE OR MORE LINES OF THERAPY
- I-724 TREATMENT OF ADULT PATIENTS WITH PROGRESSIVE, WELL DIFFERENTIATED, NON-FUNCTIONAL NEUROENDOCRINE TUMORS (NET) OF GI OR LUNG ORIGIN WITH UNRESECTABLE, LOCALLY ADVANCED OR METASTATIC DISEASE
- I-725 TREATMENT OF HORMONE RECEPTOR (HR)-POSITIVE, HER2-NEGATIVE ADVANCED OR METASTATIC BREAST CANCER IN COMBINATION THERAPY WITH PALBOCICLIB AND FULVESTRANT IN WOMEN WITH DISEASE PROGRESSION FOLLOWING ENDOCRINE THERAPY.
- I-726 EXPANSION OF THE PATIENT POPULATION TO INCLUDE PATIENTS WITH RECURRENCE OF HEPATITIS C VIRUS (HCV) GENOTYPE 1 OR 3 AFTER LIVER TRANSPLANTATION
- I-727 EXPANSION OF THE INDICATION TO INCLUDE TREATMENT OF SUBJECTS WITH GENOTYPE-1 CHRONIC HEPATITIS C VIRUS INFECTION, INCLUDING SUBJECTS WHO ARE CO-INFECTED WITH THE HUMAN IMMUNODEFICIENCY VIRUS (HIV-1) BASED ON THE RESULTS FROM THE ALLY-2 CLINICAL TRIAL
- I-728 EXPANDED INDICATION FOR USE IN ULTRASONOGRAPHY OF THE LIVER FOR CHARACTERIZATION OF FOCAL LIVER LESIONS IN ADULT AND PEDIATRIC PATIENTS
- I-729 PROVIDES FOR THE FRONTLINE INDICATION FOR THE TREATMENT OF CHRONIC LYMPHOCYTIC LEUKEMIA
- I-730 NEW INDICATION FOR THE TREATMENT OF PATIENTS WITH METASTATIC, SQUAMOUS, NON-SMALL CELL LUNG CANCER PROGRESSING AFTER PLATINUM-BASED CHEMOTHERAPY
- I-731 FOR USE IN MAGNETIC RESONANCE ANGIOGRAPHY IN ADULT AND PEDIATRIC PATIENTS (INCLUDING TERM NEONATES) TO EVALUATE KNOWN OR SUSPECTED SUPRA-AORTIC OR RENAL ARTERY DISEASE
- I-732 TREATMENT OF PEDIATRIC PATIENTS 7 TO 17 YEARS OF AGE WITH HOMOZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA TO REDUCE LDL-C, TOTAL C, NONHDL-C AND APOB AS AN ADJUNCT TO DIET, EITHER ALONE OR WITH OTHER LIPID-LOWERING TREATMENTS
- I-733 USE OF CANAGLIFLOZIN FOR INITIAL THERAPY IN COMBINATION WITH METFORMIN
- I-734 EXPANDED INDICATION FOR THE USE OF LENVIMA IN COMBINATION WITH EVEROLIMUS FOR THE TREATMENT OF PATIENTS WITH ADVANCED RCC FOLLOWING ONE PRIOR ANTI-ANGIOGENIC THERAPY.
- I-735 AS AN ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS WHEN TREATMENT WITH BOTH CANAGLIFLOZIN AND METFORMIN IS APPROPRIATE
- I-736 REVISED INDICATION TO INCLUDE THE TREATMENT OF PATIENTS WITH SMALL LYMPHOCYTIC LEUKEMIA (SLL)
- I-737 REVISED INDICATION TO INCLUDE THE TREATMENT OF PATIENTS WITH SMALL LYMPHOCYTIC LEUKEMIA (SLL) WITH 17P DELETION
- I-738 REVISIONS TO THE INDICATIONS AND USAGE SECTION WITH RESPECT TO COMPLICATED INTRA-ABDOMINAL INFECTIONS
- I-739 TO REDUCE THE RISK OF CARDIOVASCULAR DEATH IN ADULT PATIENTS WITH TYPE 2 DIABETES MELLITUS AND ESTABLISHED CARDIOVASCULAR DISEASE
- I-740 EXPANDED INDICATION FOR THE TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGE 6 YEARS AND OLDER TO INCLUDE THE G1244E, G1349D, G178R, G551S, S1251N, S1255P, S549N, OR S549R MUTATION IN THE CFTR GENE
- I-741 TREATMENT OF PATIENTS WITH MARGINAL ZONE LYMPHOMA (MZL) WHO REQUIRE SYSTEMIC THERAPY AND HAVE RECEIVED AT LEAST ONE PRIOR ANTI-CD20-BASED THERAPY
- I-742 TREATMENT OF NODAL MARGINAL ZONE LYMPHOMA
- I-743 INFORMATION ADDED TO THE LABELING FOR THE ADDITION OF THE TREATMENT OF CHRONIC HEPATITIS C VIRUS (HCV) GENOTYPE 4 (GT4) INFECTED PATIENTS WITH COMPENSATED CIRRHOSIS BASED ON RESULTS FROM STUDY M11-665
- I-744 TREATMENT OF PATIENTS WITH HEPATOCELLULAR CARCINOMA (HCC) WHO HAVE BEEN PREVIOUSLY TREATED WITH SORAFENIB

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

- I-745 MEKINIST, IN COMBINATION WITH DABRAFENIB, FOR THE TREATMENT OF PATIENTS WITH METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) WITH BRAF V600E MUTATION AS DETECTED BY AN FDA-APPROVED TEST.
- I-746 NEW INDICATION OF MAINTENANCE MONOTHERAPY TREATMENT OF BIPOLAR I DISORDER IN ADULTS
- I-747 FOR REDUCING THE RISK OF GRAFT REJECTION WHEN USED WITH HIGH-DOSE BUSULFAN AND CYCLOPHOSPHAMIDE AS A PREPARATIVE REGIMEN FOR ALLOGENEIC HEMATOPOIETIC PROGENITOR (STEM) CELL TRANSPLANTATION FOR PEDIATRIC PATIENTS WITH CLASS 3 BETA-THALASSEMIA
- I-748 TO REDUCE THE ACUTE COMPLICATIONS OF SICKLE CELL DISEASE IN ADULT AND PEDIATRIC PATIENTS FIVE YEARS OF AGE AND OLDER
- I-749 MONOTHERAPY FOR THE TREATMENT OF HORMONE RECEPTOR (HR) POSITIVE, HUMAN EPIDERMAL GROWTH FACTOR RECEPTOR 2 (HER2)-NEGATIVE, ADVANCED BREAST CANCER IN POSTMENOPAUSAL WOMEN NOT PREVIOUSLY TREATED WITH ENDOCRINE THERAPY
- I-750 REDUCE THE RISK OF MAJOR ADVERSE CARDIOVASCULAR EVENTS IN ADULTS WITH TYPE 2 DIABETES MELLITUS AND ESTABLISHED CARDIOVASCULAR DISEASE
- I-751 TREATMENT OF TARDIVE DYSKINESIA
- I-752 CORONARY COMPUTED TOMOGRAPHY ANGIOGRAPHY (CCTA) TO ASSIST DIAGNOSTIC EVALUATION OF PATIENTS WITH SUSPECTED CORONARY ARTERY DISEASE
- I-753 TREATMENT OF ADULT PATIENTS WITH CHRONIC GRAFT VERSUS HOST DISEASE (CGVHD) AFTER FAILURE OF ONE OR MORE LINES OF SYSTEMIC THERAPY
- I-754 TO REDUCE THE FREQUENCY OF SHORT-ACTING SOMATOSTATIN ANALOG RESCUE THERAPY WHEN USED FOR THE TREATMENT OF ADULTS WITH CARCINOID SYNDROME
- I-755 ADJUVANT TREATMENT OF ADULT PATIENTS AT HIGH RISK OF RECURRENT RENAL CELL CARCINOMA (RCC) FOLLOWING NEPHRECTOMY
- I-756 EXPANDED THE APPROVED INDICATION BY REMOVING THE RESTRICTION FOR USE ONLY IN PATIENTS WHO HAVE PROGRESSED ON OR ARE INTOLERANT TO CRIZOTINIB
- I-757 TREATMENT OF PATIENTS WITH ERDHEIM-CHESTER DISEASE WITH BRAF V600 MUTATION
- I-758 FOR USE WITH RILPIVIRINE AS A COMPLETE REGIMEN TO REPLACE THE CURRENT ARV REGIMEN IN VIROLOGICALLY SUPPRESSED PATIENTS ON A STABLE ARV REGIMEN FOR AT LEAST 6 MONTHS WITH NO HISTORY OF TX FAILURE OR KNOWN SUBSTITUTIONS ASSOC. WITH RESISTANCE TO EITHER ARV
- I-759 TREATMENT OF ADULT PATIENTS WITH NEWLY-DIAGNOSED CHRONIC PHASE (CP) PHILADELPHIA CHROMOSOME-POSITIVE CHRONIC MYELOGENOUS LEUKEMIA (PH+CML)
- I-760 FOR THE TREATMENT OF PATIENTS WITH ADVANCED RENAL CELL CARCINOMA
- I-761 TREATMENT OF ADULT PATIENTS WITH ACTIVE PSORIATIC ARTHRITIS WHO HAVE HAD AN INADEQUATE RESPONSE OR INTOLERANCE TO METHOTREXATE OR OTHER NON-BIOLOGIC DISEASE-MODIFYING ANTIRHEUMATIC DRUGS
- I-762 TREATMENT OF DELETERIOUS OR SUSPECTED DELETERIOUS GERMLINE BRCA-MUTATED, HUMAN EPIDERMAL GROWTH FACTOR RECEPTOR 2-NEGATIVE METASTATIC BREAST CANCER WHO HAVE BEEN TREATED WITH CHEMOTHERAPY IN THE NEOADJUVANT, ADJUVANT OR METASTATIC SETTING
- I-763 TREATMENT OF PATIENTS WITH METASTATIC NON-SMALL CELL LUNG CANCER WHOSE TUMORS HAVE NON-RESISTANT EPIDERMAL GROWTH FACTOR RECEPTOR MUTATIONS AS DETECTED BY AN FDA-APPROVED TEST
- I-764 TREATMENT IN ADULT PATIENTS FOR IRRITABLE BOWEL SYNDROME WITH CONSTIPATION (IBS-C)
- I-765 ABIRATERONE ACETATE IN COMBINATION WITH PREDNISONE FOR THE TREATMENT OF PATIENTS WITH METASTATIC HIGH-RISK CASTRATION-SENSITIVE PROSTATE CANCER
- I-766 TREATMENT OF MINIMALLY TO MODERATELY THICK ACTINIC KERATOSIS OF THE UPPER EXTREMITIES IN CONJUNCTION WITH A BLUE LIGHT PHOTODYNAMIC THERAPY ILLUMINATOR
- I-767 TREATMENT OF IRON DEFICIENCY ANEMIA IN ADULT PATIENTS WHO HAVE INTOLERANCE TO ORAL IRON OR HAVE HAD UNSATISFACTORY RESPONSE TO ORAL IRON
- I-768 IN COMBINATION WITH AN AROMATASE INHIBITOR AS INITIAL ENDOCRINE-BASED THERAPY FOR THE TREATMENT OF POSTMENOPAUSAL WOMEN WITH HORMONE RECEPTOR-POSITIVE, HUMAN EPIDERMAL GROWTH FACTOR RECEPTOR 2-NEGATIVE ADVANCED OR METASTATIC BREAST CANCER
- I-769 TREATMENT OF DYSKINESIA IN PATIENTS WITH PARKINSON'S DISEASE RECEIVING LEVODOPA-BASED THERAPY, WITH OR WITHOUT CONCOMITANT DOPAMINERGIC MEDICATIONS

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

- I-770 TREATMENT OF ACUTE OTITIS EXTERNA IN PATIENTS 6 MONTHS OF AGE AND OLDER DUE TO PSEUDOMONAS AERUGINOSA AND STAPHYLOCOCCUS AUREUS
- I-771 REVISION OF THE INDICATION SECTION OF THE PACKAGE INSERT REGARDING AN INTERSCALENE BRACHIAL PLEXUS NERVE BLOCK TO PRODUCE POSTSURGICAL REGIONAL ANALGESIA
- I-772 FOR THE MAINTENANCE TREATMENT OF ADULT PATIENTS WITH RECURRENT EPITHELIAL OVARIAN, FALLOPIAN TUBE, OR PRIMARY PERITONEAL CANCER WHO ARE IN A COMPLETE OR PARTIAL RESPONSE TO PLATINUM-BASED CHEMOTHERAPY
- I-773 FOR THE ADJUNCTIVE TREATMENT OF ADULT AND PEDIATRIC PATIENTS AGE 2 YEARS AND OLDER WITH TUBEROUS SCLEROSIS COMPLEX (TSC)-ASSOCIATED PARTIAL-ONSET SEIZURES
- I-774 TO ALLOW FOR FIRST-LINE TREATMENT OF PATIENTS WITH METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) WHOSE TUMORS HAVE EPIDERMAL GROWTH FACTOR RECEPTOR (EGFR) EXON 19 DELETIONS OR EXON 21 (L858R) SUBSTITUTION MUTATIONS, AS DETECTED BY AN FDA APPROVED TEST
- I-775 REVISED INDICATION FOR FIXED-DOSE COMBINATION OF FLUTICASONE FUROATE, UMECLIDINIUM, AND VILANTEROL TO TREAT AIRFLOW OBSTRUCTION IN CHRONIC OBSTRUCTIVE PULMONARY DISEASE (COPD) AND TO REDUCE COPD EXACERBATIONS IN PTS WITH HISTORY OF EXACERBATIONS
- I-776 FIRSTLINE MAINTENANCE TX IN PTS W/ DELETERIOUS OR SUSPECTED DELETERIOUS GERMLINE, SOMATIC BRCA-MUTATED ADVANCED EPITHELIAL OVARIAN, FALLOPIAN TUBE OR PRIMARY PERITONEAL CA WHO ARE IN COMPLETE OR PARTIAL RESPONSE TO FIRSTLINE PLATINUM-BASED CHEMOTHERAPY
- I-777 CO-ADMINISTRATION THERAPY OF MIRABEGRON WITH SOLIFENACIN SUCCINATE FOR TREATMENT OF OVERACTIVE BLADDER WITH SYMPTOMS OF URGE URINARY INCONTINENCE, URGENCY, AND URINARY FREQUENCY
- I-778 DABRAFENIB IN COMBINATION WITH TRAMETINIB FOR THE ADJUVANT TREATMENT OF PATIENTS WITH MELANOMA WITH BRAF V600E OR V600K MUTATIONS, AS DETECTED BY AN FDA-APPROVED TEST, AND INVOLVEMENT OF LYMPH NODE(S), FOLLOWING COMPLETE RESECTION
- I-779 USE OF TOLVAPTAN TO SLOW KIDNEY FUNCTION DECLINE IN ADULTS AT RISK OF RAPIDLY PROGRESSING AUTOSOMAL DOMINANT POLYCYSTIC KIDNEY DISEASE (ADPKD)
- I-780 TREATMENT OF ADULT PATIENTS WITH MODERATELY TO SEVERELY ACTIVE ULCERATIVE COLITIS (UC)
- I-781 DABRAFENIB IN COMBINATION WITH TRAMETINIB FOR THE TREATMENT OF PATIENTS WITH LOCALLY ADVANCED OR METASTATIC ANAPLASTIC THYROID CANCER (ATC) WITH BRAF V600E MUTATION AND WITH NO SATISFACTORY LOCOREGIONAL TREATMENT OPTIONS
- I-782 REVISIONS TO INDICATION FOR THE TREATMENT OF PATIENTS WITH CHRONIC LYMPHOCYTIC LEUKEMIA (CLL) OR SMALL LYMPHOCYTIC LYMPHOMA (SLL), WITH OR WITHOUT 17P DELETION, WHO HAVE RECEIVED AT LEAST ONE PRIOR THERAPY
- I-783 EXPANDED INDICATION TO INCLUDE RIBOCICLIB WITH AN AROMATASE INHIBITOR IN PRE/PERIMENOPAUSAL WOMEN WITH HORMONE RECEPTOR-POSITIVE, HUMAN EPIDERMAL GROWTH FACTOR RECEPTOR 2-NEGATIVE ADVANCED OR METASTATIC BREAST CANCER, AS INITIAL ENDOCRINE-BASED THERAPY
- I-784 RIBOCICLIB WITH FULVESTRANT FOR THE TREATMENT OF POSTMENOPAUSAL WOMEN WITH HR-POSITIVE, HER2-NEGATIVE ADVANCED OR METASTATIC BREAST CANCER, AS INITIAL ENDOCRINE BASED THERAPY OR FOLLOWING DISEASE PROGRESSION ON ENDOCRINE THERAPY
- I-785 TREATMENT OF PATIENTS WITH CUSHING'S DISEASE FOR WHOM PITUITARY SURGERY IS NOT AN OPTION OR HAS NOT BEEN CURATIVE
- I-786 TREATMENT OF PATIENTS WITH NON-METASTATIC CASTRATION-RESISTANT PROSTATE CANCER
- I-787 FIRST-LINE TREATMENT OF PATIENTS WITH UNRESECTABLE HEPATOCELLULAR CARCINOMA (HCC)
- I-788 NEW INDICATION FOR CANAGLIFLOZIN TO REDUCE THE RISK OF MAJOR ADVERSE CARDIOVASCULAR EVENTS (CARDIOVASCULAR DEATH, NONFATAL MYOCARDIAL INFARCTION AND NONFATAL STROKE) IN ADULTS WITH TYPE 2 DIABETES MELLITUS AND ESTABLISHED CARDIOVASCULAR DISEASE (CVD)
- I-789 VENETOCLAX IN COMBO WITH AZACITIDINE OR DECITABINE OR LOW-DOSE CYTARABINE FOR THE TX OF NEWLY-DIAGNOSED ACUTE MYELOID LEUKEMIA IN ADULTS WHO ARE AGE 75 YEARS OR OLDER, OR WHO HAVE COMORBIDITIES THAT PRECLUDE USE OF INTENSIVE INDUCTION CHEMOTHERAPY
- I-790 USE OF FERRIC CITRATE FOR THE TREATMENT OF IRON DEFICIENCY ANEMIA IN ADULT PATIENTS WITH CKD NOT ON DIALYSIS

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

I-791	TREATMENT OF PEDIATRIC PATIENTS ONE YEAR OF AGE AND OLDER WITH NEWLY DIAGNOSED PHILADELPHIA CHROMOSOME-POSITIVE (PH+) ACUTE LYMPHOBLASTIC LEUKEMIA (ALL) IN COMBINATION WITH CHEMOTHERAPY
I-792	TREATMENT OF PATIENTS WITH HEPATOCELLULAR CARCINOMA WHO HAVE BEEN PREVIOUSLY TREATED WITH SORAFENIB
I-793	TREATMENT OF MODERATE TO SEVERE VAGINAL DRYNESS, A SYMPTOM OF VULVAR AND VAGINAL ATROPHY, DUE TO MENOPAUSE
I-794	TREATMENT OF ADULT PATIENTS WITH METASTATIC GASTRIC OR GASTROESOPHAGEAL JUNCTION ADENOCARCINOMA PREVIOUSLY TREATED WITH AT LEAST TWO PRIOR LINES OF CHEMOTHERAPY, AND IF APPROPRIATE, HER2/NEU-TARGETED THERAPY
I-795	VENETOCLAX IN COMBINATION WITH OBINUTUZUMAB IN PREVIOUSLY UNTREATED PATIENTS WITH CHRONIC LYMPHOCYTIC LEUKEMIA OR SMALL LYMPHOCYTIC LYMPHOMA
I-796	USED IN COMBINATION WITH A RITUXIMAB PRODUCT, ARE INDICATED FOR THE TREATMENT OF ADULT PATIENTS WITH PREVIOUSLY TREATED FOLLICULAR LYMPHOMA (FL)
I-797	USED IN COMBINATION WITH A RITUXIMAB PRODUCT, ARE INDICATED FOR THE TREATMENT OF ADULT PATIENTS WITH PREVIOUSLY TREATED MARGINAL ZONE LYMPHOMA (MZL)
I-798	TREATMENT OF DEPRESSIVE EPISODES ASSOCIATED WITH BIPOLAR I DISORDER (BIPOLAR DEPRESSION)
I-799	TREATMENT OF STEROID-REFRACTORY ACUTE GRAFT-VERSUS-HOST DISEASE (GVHD) IN ADULT AND PEDIATRIC PATIENTS 12 YEARS AND OLDER
I-800	TREATMENT OF OCULAR INFLAMMATION FOLLOWING OPHTHALMIC SURGERY
I-801	USE IN CARDIAC MRI TO ASSESS MYOCARDIAL PERFUSION (STRESS, REST) AND LATE GADOLINIUM ENHANCEMENT IN ADULT PATIENTS WITH KNOWN OR SUSPECTED CORONARY ARTERY DISEASE (CAD)
I-802	TREATMENT OF THROMBOCYTOPENIA IN ADULT PATIENTS WITH CHRONIC IMMUNE THROMBOCYTOPENIA WHO HAVE HAD AN INSUFFICIENT RESPONSE TO A PREVIOUS TREATMENT
I-803	TREATMENT OF ADULT PATIENTS WITH ORAL ULCERS ASSOCIATED WITH BEHCETS DISEASE
I-804	EXPANDED INDICATION FOR USE AS AN ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS
I-805	SLOW THE RATE OF DECLINE IN PULMONARY FUNCTION IN PATIENTS WITH SYSTEMIC SCLEROSIS-ASSOCIATED INTERSTITIAL LUNG DISEASE
I-806	EXPANDED INDICATION FOR PTS WHO ARE VIROLOGICALLY SUPPRESSED (HIV-1 RNA LESS THAN 50 COPIES/ML) ON A STABLE ARV REGIMEN WITH NO HX OF TX FAILURE AND NO KNOWN SUBSTITUTIONS ASSOCIATED W RESISTANCE TO DORAVIRINE, LAMIVUDINE OR TENOFOVIR DISOPROXIL FUMARATE
I-807	TREATMENT OF ADVANCED ENDOMETRIAL CARCINOMA THAT IS NOT MICROSATELLITE INSTABILITY-HIGH OR MISMATCH REPAIR DEFICIENT, WHO HAVE DISEASE PROGRESSION FOLLOWING PRIOR SYSTEMIC THERAPY AND ARE NOT CANDIDATES FOR CURATIVE SURGERY OR RADIATION
I-808	TREATMENT OF PATIENTS WITH METASTATIC CASTRATION-SENSITIVE PROSTATE CANCER (MCSPC)
I-809	TO REDUCE THE RISK OF END-STAGE KIDNEY DISEASE, DOUBLING OF SERUM CREATININE, CARDIOVASCULAR DEATH, AND HOSPITALIZATION FOR HEART FAILURE IN ADULTS WITH TYPE 2 DIABETES MELLITUS AND DIABETIC NEPHROPATHY WITH ALBUMINURIA > 300 MG/DAY
I-810	PROPHYLAXIS OF VENOUS THROMBOEMBOLISM IN ACUTELY ILL MEDICAL PATIENTS AT RISK FOR THROMBOEMBOLIC COMPLICATIONS NOT AT HIGH RISK OF BLEEDING
I-811	TREATMENT OF ACUTE UNCOMPLICATED INFLUENZA IN PATIENTS 12 YEARS OF AGE OR OLDER, WHO HAVE BEEN SYMPTOMATIC FOR NO MORE THAN 48 HOURS AND ARE AT HIGH RISK OF DEVELOPING INFLUENZA-RELATED COMPLICATIONS
I-812	FOR USE IN AT RISK ADULTS AND ADOLESCENTS WEIGHING AT LEAST 35 KG FOR PRE-EXPOSURE PROPHYLAXIS TO REDUCE THE RISK OF HIV-1 INFECTION FROM SEXUAL ACQUISITION, EXCLUDING INDIVIDUALS AT RISK FROM RECEPTIVE VAGINAL SEX
I-813	TX OF ADULT PTS W/ ADV OVARIAN FALLOPIAN TUBE OR PRIMARY PERITONEAL CANCER TREATED W/ >=3 PRIOR CHEMO REGIMENS & ASSOCIATED W/ HRD DEFICIENCY POSITIVE STATUS DEFINED BY A DELETERIOUS OR SUSPECTED DELETERIOUS BRCA MUTATION
I-814	TX OF ADV OVARIAN FALLOPIAN TUBE OR PRIMARY PERITONEAL CANCER TREATED W/ >=3 PRIOR CHEMO REGIMENS & ASSOCIATED W/ HRD DEFICIENCY DEFINED BY POSITIVE STATUS GENOMIC INSTABILITY & WHO HAVE PROGRESSED >6MO AFTER RESPONSE TO LAST PLATINUM-BASED CHEMO

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

- I-815 TREATMENT OF COMMUNITY ACQUIRED BACTERIAL PNEUMONIA (CABP) CAUSED BY DESIGNATED SUSCEPTIBLE BACTERIA IN ADULTS
- I-816 TREATMENT OF ADULT PATIENTS WITH NEWLY-DIAGNOSED ACUTE MYELOID LEUKEMIA (AML) WHO ARE \geq 75 YEARS OLD OR WHO HAVE COMORBIDITIES THAT PRECLUDE USE OF INTENSIVE INDUCTION CHEMOTHERAPY
- I-817 TREATMENT OF ADULT PATIENTS WITH CHRONIC LYMPHOCYTIC LEUKEMIA (CLL) OR SMALL LYMPHOCYTIC LYMPHOMA (SLL)
- I-818 MAINTENANCE TREATMENT OF ADULT PATIENTS WITH DELETERIOUS OR SUSPECTED DELETERIOUS GBRCA METASTATIC PANCREATIC ADENOCARCINOMA WHOSE DISEASE HAS NOT PROGRESSED ON AT LEAST 16 WEEKS OF A FIRST-LINE PLATINUM-BASED CHEMOTHERAPY REGIMEN
- I-819 ADJUNCT TO MAX TOLERATED STATIN TX TO REDUCE RISK OF MI, STROKE, CORONARY REVASCULARIZATION, & UNSTABLE ANGINA REQUIRING HOSPITALIZATION IN ADULTS W/ ELEVATED TG LEVELS & ESTABLISHED CV DISEASE OR DIABETES MELLITUS & 2+ RISK FACTORS FOR CV DISEASE
- I-820 INDICATED FOR THE TREATMENT OF PULMONARY ARTERIAL HYPERTENSION (PAH) (WHO GROUP 1) TO DELAY DISEASE PROGRESSION
- I-821 TREATMENT OF CANDIDEMIA, ACUTE DISSEMINATED CANDIDIASIS, CANDIDA PERITONITIS AND ABSCESSSES WITHOUT MENINGOENCEPHALITIS AND/OR OCULAR DISSEMINATION IN PEDIATRIC PATIENTS YOUNGER THAN 4 MONTHS OF AGE
- I-822 REDUCE THE RISK OF MAJOR ADVERSE CARDIOVASCULAR EVENTS (CARDIOVASCULAR DEATH, NON-FATAL MYOCARDIAL INFARCTION OR NON-FATAL STROKE) IN ADULTS WITH TYPE 2 DIABETES MELLITUS AND ESTABLISHED CARDIOVASCULAR DISEASE
- I-823 USE IN COMBINATION WITH CAPECITABINE, FOR THE TREATMENT OF ADULT PATIENTS WITH ADVANCED OR METASTATIC HER2-POSITIVE BREAST CANCER WHO HAVE RECEIVED TWO OR MORE PRIOR ANTI-HER2 BASED REGIMENS IN THE METASTATIC SETTING
- I-824 RIVAROXABAN IN COMBINATION WITH ASPIRIN, IS INDICATED TO REDUCE THE RISK OF MAJOR CV EVENTS (CV DEATH, MI, AND STROKE) IN PATIENTS WITH CHRONIC CORONARY ARTERY DISEASE (CAD) OR PERIPHERAL ARTERY DISEASE (PAD)
- I-825 TREATMENT FOR CHRONIC FIBROSING INTERSTITIAL LUNG DISEASES WITH A PROGRESSIVE PHENOTYPE
- I-826 ENCORAFENIB, IN COMBINATION WITH CETUXIMAB, FOR THE TREATMENT OF ADULT PATIENTS WITH METASTATIC COLORECTAL CANCER WITH A BRAF V600E MUTATION, AS DETECTED BY AN FDA-APPROVED TEST, AFTER PRIOR THERAPY
- I-827 EXPANDED INDICATION FOR PATIENTS WHO ARE VIROLOGICALLY SUPPRESSED (HIV-1 RNA LESS THAN 50 COPIES/ML) ON A STABLE ARV REGIMEN WITH NO HISTORY OF TREATMENT FAILURE AND NO KNOWN SUBSTITUTIONS ASSOCIATED WITH RESISTANCE TO DORAVIRINE
- I-828 INDICATED FOR THE TREATMENT OF ADULT PATIENTS WITH CUSHING'S DISEASE FOR WHOM PITUITARY SURGERY IS NOT AN OPTION OR HAS NOT BEEN CURATIVE
- I-829 TREATMENT OF PEDIATRIC PATIENTS 2 YEARS OF AGE AND OLDER WITH CENTRAL PRECOCIOUS PUBERTY (CPP)
- I-830 TREATMENT OF ADULT PATIENTS WITH A DELETERIOUS BRCA MUTATION (GERMLINE AND/OR SOMATIC)-ASSOCIATED METASTATIC CASTRATION-RESISTANT PROSTATE CANCER WHO HAVE BEEN TREATED WITH ANDROGEN RECEPTOR-DIRECTED THERAPY AND A TAXANE-BASED CHEMOTHERAPY
- I-831 W/BEVACIZUMAB FOR MAINTENANCE TX OF ADULTS W/ADV. EPITHELIAL OVARIAN, FALLOPIAN TUBE OR PRIMARY PERITONEAL CA IN COMPLETE OR PARTIAL RESPONSE TO FIRST-LINE PLATINUM-BASED CHEMO & CA ASSOCIATED W/ HOMOLOGOUS RECOMBINATION DEFICIENCY POSITIVE STATUS
- I-832 TX OF ADULT PTS WITH DELETERIOUS OR SUSPECTED DELETERIOUS GERMLINE OR SOMATIC HOMOLOGOUS RECOMBINATION REPAIR GENE-MUTATED METASTATIC CASTRATION-RESISTANT PROSTATE CANCER WHO HAVE PROGRESSED FOLLOWING PRIOR TREATMENT WITH ENZALUTAMIDE OR ABIRATERONE
- I-833 MAINTENANCE TREATMENT OF ADULT PATIENTS WITH ADVANCED EPITHELIAL OVARIAN, FALLOPIAN TUBE, OR PRIMARY PERITONEAL CANCER WHO ARE IN A COMPLETE OR PARTIAL RESPONSE TO FIRST-LINE PLATINUM-BASED CHEMOTHERAPY
- I-834 TO REDUCE THE RISK OF CARDIOVASCULAR DEATH AND HOSPITALIZATION FOR HEART FAILURE IN ADULTS WITH HEART FAILURE (NYHA CLASS II-IV) WITH REDUCED EJECTION FRACTION
- I-835 TREATMENT OF ADULT PATIENTS WITH RELAPSED OR REFRACTORY (R/R) FOLLICULAR LYMPHOMA (FL) WHOSE TUMORS ARE POSITIVE FOR AN EZH2 MUTATION AS DETECTED BY AN FDA-APPROVED TEST AND WHO HAVE RECEIVED AT LEAST 2 PRIOR SYSTEMIC THERAPIES

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

- I-836 TREATMENT OF ADULT PATIENTS WITH RELAPSED OR REFRACTORY (R/R) FOLLICULAR LYMPHOMA (FL) WHO HAVE NO SATISFACTORY ALTERNATIVE TREATMENT OPTIONS
- I-837 TREATMENT OF ADULT PATIENTS WITH RELAPSED OR REFRACTORY DIFFUSE LARGE B-CELL LYMPHOMA (DLBCL), NOT OTHERWISE SPECIFIED, INCLUDING DLBCL ARISING FROM FOLLICULAR LYMPHOMA, AFTER AT LEAST 2 LINES OF SYSTEMIC THERAPY
- I-838 TREATMENT OF NEUROPATHIC PAIN ASSOCIATED WITH DIABETIC PERIPHERAL NEUROPATHY (DPN) OF THE FEET
- I-839 TO REPLACE THE CURRENT ANTIRETROVIRAL REGIMEN IN THOSE WHO ARE VIROLOGICALLY SUPRESSED (HIV-1 RNA LESS THAN 50 COPIES PER ML) ON A STABLE ANTIRETROVIRAL REGIMEN WITH NO HISTORY OF TREATMENT FAILURE
- I-840 TREATMENT OF SYMPTOMS IN ADULTS WITH MAJOR DEPRESSIVE DISORDER (MDD) WITH ACUTE SUICIDAL IDEATION OR BEHAVIOR.
- I-841 TO REDUCE THE RISK OF HOSPITALIZATION FOR HEART FAILURE IN ADULTS WITH TYPE 2 DIABETES MELLITUS AND ESTABLISHED CARDIOVASCULAR DISEASE OR MULTIPLE CARDIOVASCULAR RISK FACTORS
- I-842 TREATMENT OF ADULT PATIENTS WITH RELAPSED OR REFRACTORY MULTIPLE MYELOMA WHO HAVE RECEIVED ONE TO THREE LINES OF THERAPY IN COMBINATION WITH DARATUMUMAB AND DEXAMETHASONE
- I-843 MAINTENANCE TREATMENT OF ASTHMA IN PATIENTS AGED 18 YEARS AND OLDER
- I-844 INDICATED IN PATIENTS 18 YEARS OF AGE AND OLDER FOR THE TREATMENT OF HOSPITAL ACQUIRED BACTERIAL PNEUMONIA AND VENTILATOR-ASSOCIATED BACTERIAL PNEUMONIA (HABP/VABP) CAUSED BY THE FOLLOWING SUSCEPTIBLE GRAM-NEGATIVE MICROORGANISMS: ACINETOBACTER BAUMANNII COMPLEX, ESCHERICHIA COLI, ENTEROBACTER CLOACAE COMPLEX, KLEBSIELLA PNEUMONIAE, PSEUDOMONAS AERUGINOSA, AND SERRATIA MARCESCENS
- I-845 TREATMENT OF ADULT PATIENTS WITH ANAPLASTIC LYMPHOMA KINASE (ALK)-POSITIVE METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) AS DETECTED BY AN FDA-APPROVED TEST
- I-846 TREATMENT OF CATAPLEXY IN ADULT PATIENTS WITH NARCOLEPSY
- I-847 EXPANDED INDICATION OF THE TREATMENT OF ADULT PATIENTS WITH ANAPLASTIC LYMPHOMA KINASE (ALK)-POSITIVE METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) AS DETECTED BY AN FDA-APPROVED TEST
- I-848 REDUCE THE RISK OF STROKE IN PATIENTS WITH ACUTE ISCHEMIC STROKE (NIH STROKE SCALE SCORE \leq 5) OR HIGH-RISK TRANSIENT ISCHEMIC ATTACK (TIA)
- I-849 CHRONIC PHASE (CP) CHRONIC MYELOID LEUKEMIA (CML) WITH RESISTANCE OR INTOLERANCE TO AT LEAST TWO PRIOR KINASE INHIBITORS
- I-850 TREATMENT OF NIGHTTIME SLEEP DISTURBANCES IN SMITH-MAGENIS SYNDROME (SMS) IN PATIENTS 16 YEARS OF AGE AND OLDER
- I-851 TO REDUCE THE RISK OF A FIRST MYOCARDIAL INFARCTION (MI) OR STROKE IN PATIENTS WITH CORONARY ARTERY DISEASE (CAD) AT HIGH RISK FOR SUCH EVENTS
- I-852 TREATMENT OF PEDIATRIC PATIENTS 1 YEAR OF AGE AND OLDER AND YOUNG ADULTS WITH RELAPSED OR REFRACTORY, SYSTEMIC ANAPLASTIC LARGE CELL LYMPHOMA (ALCL) THAT IS ALK-POSITIVE
- I-853 INDICATION OF OSIMERTINIB AS ADJUVANT THERAPY AFTER TUMOR RESECTION IN ADULT PATIENTS WITH NON-SMALL CELL LUNG CANCER (NSCLC) WHOSE TUMORS HAVE EPIDERMAL GROWTH FACTOR RECEPTOR (EGFR) EXON 19 DELETIONS OR EXON 21 L858R MUTATIONS, AS DETECTED BY AN FDA-APPROVED TEST
- I-854 FOR THE TREATMENT OF PATIENTS WITH ADVANCED RENAL CELL CARCINOMA, AS A FIRST-LINE TREATMENT IN COMBINATION WITH NIVOLUMAB
- I-855 TREATMENT OF NEUROGENIC DETRUSOR OVERACTIVITY (NDO) IN PEDIATRIC PATIENTS 3 YEARS AND OLDER AND WEIGHING 35 KILOGRAMS OR MORE
- I-856 INDICATION FOR THE TREATMENT OF PULMONARY HYPERTENSION ASSOCIATED WITH INTERSTITIAL LUNG DISEASE TO IMPROVE EXERCISE ABILITY
- I-857 TO REDUCE THE RISK OF SUSTAINED EGFR DECLINE, END-STAGE KIDNEY DISEASE, CARDIOVASCULAR DEATH, AND HOSPITALIZATION FOR HEART FAILURE IN ADULTS WITH CHRONIC KIDNEY DISEASE AT RISK OF PROGRESSION
- I-858 FOR THE TOPICAL TREATMENT OF SCABIES INFESTATIONS IN ADULT AND PEDIATRIC PATIENTS 4 YEARS OF AGE AND OLDER
- I-859 TREATMENT OF PATIENTS WITH TRANSFUSIONAL IRON OVERLOAD DUE TO SICKLE CELL DISEASE OR OTHER ANEMIAS

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

- I-860 FOR THE TREATMENT OF MODERATELY TO SEVERELY ACTIVE ULCERATIVE COLITIS IN ADULT PATIENTS
- I-861 TREATMENT OF NEUROGENIC DETRUSOR OVERACTIVITY (NDO) IN PEDIATRIC PATIENTS 6 YEARS OF AGE AND OLDER AND WEIGHING GREATER THAN 25 KG
- I-862 TREATMENT OF VENOUS THROMBOEMBOLIC EVENTS (VTE) IN PEDIATRIC PATIENTS 8 TO LESS THAN 18 YEARS OF AGE WHO HAVE BEEN TREATED WITH A PARENTERAL ANTICOAGULANT FOR AT LEAST 5 DAYS AND TO REDUCE THE RISK OF RECURRENCE OF VTE IN PEDIATRIC PATIENTS 8 TO LESS THAN 18 YEARS OF AGE WHO HAVE BEEN PREVIOUSLY TREATED
- I-863 TREATMENT OF ADULT PATIENTS WITH ADVANCED SYSTEMIC MASTOCYTOSIS (ADVSM), INCLUDING PATIENTS WITH AGGRESSIVE SYSTEMIC MASTOCYTOSIS (ASM) AND SYSTEMIC MASTOCYTOSIS WITH AN ASSOCIATED HEMATOLOGICAL NEOPLASM (SM-AHN)
- I-864 TREATMENT OF ADULT PATIENTS WITH MAST CELL LEUKEMIA (MCL)
- I-865 FOR THE PREVENTIVE TREATMENT OF EPISODIC MIGRAINE IN ADULTS
- I-866 FOR THE TREATMENT OF TRICHOMONIASIS CAUSED BY TRICHOMONAS VAGINALIS IN ADULTS
- I-867 INDICATED TO REDUCE THE RISK OF MAJOR THROMBOTIC VASCULAR EVENTS (MYOCARDIAL INFARCTION, ISCHEMIC STROKE, ACUTE LIMB ISCHEMIA, AND MAJOR AMPUTATION OF VASCULAR ETIOLOGY) IN PATIENTS WITH PAD, INCLUDING PATIENTS WHO HAVE RECENTLY UNDERGONE A LOWER EXTREMITY REVASCLARIZATION PROCEDURE DUE TO SYMPTOMATIC PAD
- I-868 LENVATINIB IN COMBINATION WITH PEMBROLIZUMAB, IS INDICATED FOR THE FIRST-LINE TREATMENT OF ADULT PATIENTS WITH ADVANCED RENAL CELL CARCINOMA (RCC)
- I-869 REDUCE THE RISK OF CARDIOVASCULAR DEATH AND HOSPITALIZATION FOR HEART FAILURE IN ADULTS WITH HEART FAILURE AND REDUCED EJECTION FRACTION
- I-870 INDICATED FOR THE TREATMENT OF IDIOPATHIC HYPERSOMNIA (IH) IN ADULTS
- I-871 TREATMENT OF ADULT PATIENTS WITH WALDENSTROM'S MACROGLOBULINEMIA (WM)
- I-872 ADDITION OF THE INDICATION OF TREATMENT OF CHRONIC GRAFT-VERSUS-HOST DISEASE (CGVHD) AFTER FAILURE OF ONE OR TWO LINES OF SYSTEMIC THERAPY IN ADULT AND PEDIATRIC PATIENTS 12 YEARS AND OLDER
- I-873 TREATMENT OF ADULT AND PEDIATRIC PATIENTS 12 YEARS OF AGE AND OLDER WITH LOCALLY ADVANCED OR METASTATIC DIFFERENTIATED THYROID CANCER (DTC) THAT HAS PROGRESSED FOLLOWING PRIOR VEGFR-TARGETED THERAPY AND WHO ARE RADIOACTIVE IODINE-REFRACTORY OR INELIGIBLE
- I-874 TREATMENT OF ADULT PATIENTS WITH RELAPSED OR REFRACTORY MARGINAL ZONE LYMPHOMA (MZL) WHO HAVE RECEIVED AT LEAST ONE ANTI-CD20-BASED REGIMEN
- I-875 FOR THE TREATMENT OF ADULT PATIENTS WITH PREVIOUSLY TREATED, LOCALLY ADVANCED OR METASTATIC CHOLANGIOCARCINOMA WITH AN IDH1 MUTATION AS DETECTED BY AN FDA-APPROVED TEST
- I-876 TREATMENT OF OCULAR ITCHING ASSOCIATED WITH ALLERGIC CONJUNCTIVITIS
- I-877 INDICATION FOR THE USE OF ABEMACICLIB IN COMBINATION WITH ENDOCRINE THERAPY (TAMOXIFEN OR AN AROMATASE INHIBITOR) FOR THE ADJUVANT TREATMENT OF ADULT PATIENTS WITH HORMONE RECEPTOR (HR)-POSITIVE, HUMAN EPIDERMAL GROWTH FACTOR RECEPTOR 2 (HER2)-NEGATIVE, NODE-POSITIVE, EARLY CANCER (EBC) AT HIGH RISK OF RECURRENCE AND A KI-67 SCORE>20% AS DETERMINED BY AN FDA APPROVED TEST
- I-878 ADDITION OF A NEW INDICATION FOR ADJUNCTIVE THERAPY IN THE TREATMENT OF PRIMARY GENERALIZED TONIC-CLONIC SEIZURES IN PATIENTS 4 YEARS OF AGE AND OLDER
- I-879 TREATMENT OF ADULT PATIENTS WITH ACTIVE ANKYLOSING SPONDYLITIS WHO HAVE HAD AN INADEQUATE RESPONSE OR INTOLERANCE TO ONE OR MORE TNF BLOCKERS, TO THE PRESCRIBING INFORMATION
- I-880 TREATMENT OF ADULTS WITH ACTIVE PSORIATIC ARTHRITIS WHO HAVE HAD AN INADEQUATE RESPONSE OR INTOLERANCE TO ONE OR MORE TNF BLOCKERS
- I-881 FOR THE TREATMENT OF INVASIVE ASPERGILLOSIS IN PATIENTS 13 YEARS OF AGE AND OLDER
- I-882 INDICATED FOR DEPRESSIVE EPISODES ASSOCIATED WITH BIPOLAR I OR II DISORDER (BIPOLAR DEPRESSION) IN ADULTS, AS MONOTHERAPY AND AS ADJUNCTIVE THERAPY WITH LITHIUM OR VALPROATE
- I-883 TREATMENT OF ADULTS AND PEDIATRIC PATIENTS 12 YEARS OF AGE AND OLDER WITH REFRACTORY, MODERATE-TO-SEVERE ATOPIC DERMATITIS WHOSE DISEASE IS NOT ADEQUATELY CONTROLLED WITH OTHER SYSTEMIC DRUG PRODUCTS, INCLUDING BIOLOGICS, OR WHEN USE OF THOSE THERAPIES ARE INADVISABLE

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

- I-884 REVISIONS TO THE LABELING TO INCLUDE DATA FOR SUBJECTS WITH MILD TO MODERATE PLAQUE PSORIASIS, AND TO ALLOW FOR AN EXPANSION OF THE INDICATION
- I-885 FOR THE ADJUVANT TREATMENT OF ADULT PATIENTS WITH DELETERIOUS OR SUSPECTED DELETERIOUS GBRCA MUTATED HER2-NEGATIVE HIGH-RISK EARLY BREAST CANCER WHO HAVE PREVIOUSLY BEEN TREATED WITH NEOADJUVANT OR ADJUVANT CHEMOTHERAPY
- I-886 TREATMENT OF ADULT PATIENTS WITH MODERATELY TO SEVERELY ACTIVE ULCERATIVE COLITIS WHO HAVE HAD AN INADEQUATE RESPONSE OR INTOLERANCE TO ONE OR MORE TNF BLOCKERS
- I-887 INDICATION FOR THE TREATMENT OF SEIZURES ASSOCIATED WITH LENNOX-GASTAUT SYNDROME (LGS) IN PATIENTS WHO ARE 2 YEARS OF AGE AND OLDER
- I-888 TREATMENT OF ADULTS WITH ACTIVE ANKYLOSING SPONDYLITIS WHO HAVE HAD AN INADEQUATE RESPONSE OR INTOLERANCE TO ONE OR MORE TNF BLOCKERS
- I-889 TREATMENT OF PEDIATRIC PATIENTS AGED ONE MONTH AND OLDER WITH NEWLY DIAGNOSED JUVENILE MYELOMONOCYTIC LEUKEMIA (JMML)
- I-890 TREATMENT OF ADULT PATIENTS WITH SEVERE ALOPECIA AREATA
- I-891 TREATMENT OF COVID-19 IN HOSPITALIZED ADULTS REQUIRING SUPPLEMENTAL OXYGEN, NONINVASIVE OR INVASIVE MECHANICAL VENTILATION, OR EXTRACORPOREAL MEMBRANE OXYGENATION (ECMO)
- I-892 CHRONIC WEIGHT MANAGEMENT IN ADULT AND PEDIATRIC PATIENTS 6 YEARS OF AGE AND OLDER WITH BARDET-BIEDL SYNDROME (BBS)
- I-893 IN COMBINATION WITH AZACITIDINE OR AS MONOTHERAPY FOR THE TREATMENT OF NEWLY DIAGNOSED ACUTE MYELOID LEUKEMIA (AML) WITH A SUSCEPTIBLE ISOCITRATE DEHYDROGENASE-1 (IDH1) MUTATION AS DETECTED BY AN FDA-APPROVED TEST IN ADULTS 75 YEARS OR OLDER, OR WHO HAVE COMORBIDITIES THAT PRECLUDE USE OF INTENSIVE INDUCTION CHEMOTHERAPY
- I-894 DABRAFENIB IS INDICATED IN COMBINATION WITH TRAMETINIB, FOR THE TREATMENT OF ADULT AND PEDIATRIC PATIENTS 6 YEARS OF AGE AND OLDER WITH UNRESECTABLE OR METASTATIC SOLID TUMORS WITH BRAF V600E MUTATION WHO HAVE PROGRESSED FOLLOWING PRIOR TREATMENT AND HAVE NO SATISFACTORY ALTERNATIVE TREATMENT OPTIONS
- I-895 TRAMETINIB IS INDICATED IN COMBINATION WITH DABRAFENIB, FOR THE TREATMENT OF ADULT AND PEDIATRIC PATIENTS 6 YEARS OF AGE AND OLDER WITH UNRESECTABLE OR METASTATIC SOLID TUMORS WITH BRAF V600E MUTATION WHO HAVE PROGRESSED FOLLOWING PRIOR TREATMENT AND HAVE NO SATISFACTORY ALTERNATIVE TREATMENT OPTIONS
- I-896 INDICATED FOR THE TOPICAL TREATMENT OF NONSEGMENTAL VITILIGO IN ADULT AND PEDIATRIC PATIENTS 12 YEARS OF AGE AND OLDER
- I-897 TREATMENT OF ADULT AND PEDIATRIC PATIENTS 1 YEAR OF AGE AND OLDER WITH UNRESECTABLE, RECURRENT OR REFRACTORY INFLAMMATORY MYOFIBROBLASTIC TUMOR (IMT) THAT IS ANAPLASTIC LYMPHOMA KINASE (ALK) POSITIVE
- I-898 FOR THE MANAGEMENT OF MODERATE TO SEVERE PAIN ASSOCIATED WITH ENDOMETRIOSIS
- I-899 TREATMENT OF ADULTS WITH RELAPSED OR REFRACTORY MYELOID/LYMPHOID NEOPLASMS (MLNS) WITH FIBROBLAST GROWTH FACTOR RECEPTOR 1 (FGFR1) REARRANGEMENT
- I-900 TREATMENT OF ADULT PATIENTS WITH METASTATIC HORMONESENSITIVE PROSTATE CANCER (MHSPC) IN COMBINATION WITH DOCETAXEL
- I-901 EXPANDED INDICATION TO INCLUDE LOWERING OF PLASMA OXALATE LEVELS IN ADULT AND PEDIATRIC PATIENTS WITH PRIMARY HYPEROXALURIA TYPE 1 (PH1)
- I-902 TREATMENT OF ADULT PATIENTS WITH HISTIOCYTIC NEOPLASMS
- I-903 REDUCTION IN THE INCIDENCE OF RECURRENT VULVOVAGINAL CANDIDIASIS (RVVC) IN ADULT AND POST-MENARCHAL PEDIATRIC FEMALES
- I-904 ADJUNCTIVE THERAPY TO ANTIDEPRESSANTS FOR THE TREATMENT OF MAJOR DEPRESSIVE DISORDER IN ADULTS
- I-905 ADJUNCT FOR INTRAOPERATIVE IDENTIFICATION OF MALIGNANT AND NONMALIGNANT PULMONARY LESIONS IN ADULT PATIENTS WITH KNOWN OR SUSPECTED CANCER IN THE LUNG
- I-906 TUCATINIB IN COMBINATION WITH TRASTUZUMAB FOR THE TREATMENT OF ADULT PATIENTS WITH RAS WILD-TYPE, HER2-POSITIVE, UNRESECTABLE OR METASTATIC COLORECTAL CANCER THAT HAS PROGRESSED FOLLOWING TREATMENT WITH FLUOROPYRIMIDINE-, OXALIPLATIN-, AND IRINOTECAN-BASED CHEMOTHERAPY
- I-907 TO INCREASE BONE DENSITY IN MEN WITH OSTEOPOROSIS AT HIGH RISK FOR FRACTURE (DEFINED AS A HISTORY OF OSTEOPOROTIC FRACTURE OR MULTIPLE RISK FACTORS FOR FRACTURE), OR PATIENTS WHO HAVE FAILED OR ARE INTOLERANT TO OTHER AVAILABLE

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

OSTEOPOROSIS THERAPY

- I-908 TREATMENT OF PEDIATRIC PATIENTS 1 YEAR OF AGE AND OLDER WITH LOW-GRADE GLIOMA WITH A BRAF V600 MUTATION WHO REQUIRE SYSTEMIC THERAPY
- I-909 PREVENTIVE TREATMENT OF MIGRAINE IN ADULTS
- I-911 FOR FLUORESCENCE IMAGING OF LYMPH NODES AND DELINEATION OF LYMPHATIC VESSELS DURING LYMPHATIC MAPPING IN ADULTS WITH BREAST CANCER FOR WHICH THIS PROCEDURE IS A COMPONENT OF INTRAOPERATIVE MANAGEMENT
- I-912 TREATMENT OF ADULT PATIENTS WITH INDOLENT SYSTEMIC MASTOCYTOSIS (ISM)
- I-913 TREATMENT OF AGITATION ASSOCIATED WITH DEMENTIA DUE TO ALZHEIMER'S DISEASE
- I-914 IN COMBINATION WITH ABIRATERONE AND PREDNISONE OR PREDNISOLONE FOR THE TREATMENT OF ADULT PATIENTS WITH DELETERIOUS OR SUSPECTED DELETERIOUS BRCA-MUTATED (BRCAM) METASTATIC CASTRATION-RESISTANT PROSTATE CANCER (MCRPC)
- I-915 TREATMENT OF IRON DEFICIENCY IN ADULT PATIENTS WITH HEART FAILURE AND NEW YORK HEART ASSOCIATION CLASS II/III TO IMPROVE EXERCISE CAPACITY
- I-916 PROPHYLAXIS OF CYTOMEGALOVIRUS (CMV) DISEASE IN ADULT KIDNEY TRANSPLANT RECIPIENTS AT HIGH RISK (DONOR CMV SEROPOSITIVE/RECIPIENT CMV SERONEGATIVE [D+/R-])
- I-917 TREATMENT OF HEAVY MENSTRUAL BLEEDING FOR UP TO 5 YEARS IN PATIENTS WHO CHOOSE INTRAUTERINE CONTRACEPTION AS THEIR METHOD OF CONTRACEPTION
- I-918 TREATMENT OF CHOLESTATIC PRURITUS IN PATIENTS 12 MONTHS OF AGE AND OLDER WITH ALAGILLE SYNDROME (ALGS)
- I-919 TREATMENT OF ADULT PATIENTS WITH MODERATELY TO SEVERELY ACTIVE CROHN'S DISEASE WHO HAVE HAD AN INADEQUATE RESPONSE OR INTOLERANCE TO ONE OR MORE TNF BLOCKERS
- I-920 USE OF TALAZOPARIB IN COMBINATION WITH ENZALUTAMIDE FOR THE TREATMENT OF ADULT PATIENTS WITH HOMOLOGOUS RECOMBINATION REPAIR (HRR) GENE-MUTATED METASTATIC CASTRATION-RESISTANT PROSTATE CANCER (MCRPC) .
- I-921 TREATMENT OF FUNCTIONAL CONSTIPATION IN PEDIATRIC PATIENTS 6 TO 17 YEARS OF AGE
- I-922 USE OF EMPAGLIFLOZIN TO REDUCE THE RISK OF SUSTAINED DECLINE IN EGFR, END-STAGE KIDNEY DISEASE, CARDIOVASCULAR DEATH, AND HOSPITALIZATION IN ADULTS WITH CHRONIC KIDNEY DISEASE AT RISK OF PROGRESSION
- I-923 FOR THE TREATMENT OF PEDIATRIC PATIENTS 1 YEAR OF AGE AND OLDER WITH CHRONIC PHASE (CP) PHILADELPHIA CHROMOSOME-POSITIVE CHRONIC MYELOGENOUS LEUKEMIA (PH+ CML), NEWLY-DIAGNOSED OR RESISTANT OR INTOLERANT TO PRIOR THERAPY
- I-924 FOR THE TREATMENT OF ADULT PATIENTS WITH RELAPSED OR REFRACTORY MYELODYSPLASTIC SYNDROMES (MDS) WITH A SUSCEPTIBLE ISOCITRATE DEHYDROGENASE-1 (IDH1) MUTATION AS DETECTED BY AN FDA-APPROVED TEST
- I-925 TREATMENT OF ADULTS WITH CHOREA ASSOCIATED WITH HUNTINGTON'S DISEASE
- I-926 TREATMENT OF PATIENTS WITH NON-METASTATIC CASTRATION-SENSITIVE PROSTATE CANCER (NMCSPC) WITH BIOCHEMICAL RECURRENCE AT HIGH RISK FOR METASTASIS
- I-928 ENCORAFENIB IN COMBINATION WITH BINIMETINIB, IS INDICATED FOR THE TREATMENT OF ADULT PATIENTS WITH METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) WITH BRAF V600E MUTATION, AS DETECTED BY AN FDA-APPROVED TEST
- I-929 INDICATED TO PRODUCE POSTSURGICAL REGIONAL ANALGESIA IN ADULTS VIA A SCIATIC NERVE BLOCK IN THE POPLITEAL FOSSA AND VIA AN ADDUCTOR CANAL BLOCK
- I-930 TREATMENT OF ADULT PATIENTS WITH LOCALLY ADVANCED OR METASTATIC UROTHELIAL CARCINOMA (MUC) WITH SUSCEPTIBLE FGFR3 GENETIC ALTERATIONS WHOSE DISEASE HAS PROGRESSED ON OR AFTER AT LEAST ONE LINE OF PRIOR SYSTEMIC THERAPY
- I-931 TREATMENT OF ADULT PATIENTS WITH ADVANCED RENAL CELL CARCINOMA FOLLOWING A PROGRAMMED DEATH RECEPTOR-1 OR PROGRAMMED DEATH-LIGAND 1 INHIBITOR AND A VASCULAR ENDOTHELIAL GROWTH FACTOR TYROSINE KINASE INHIBITOR
- I-932 IN COMBINATION WITH OXALIPLATIN, FLUOROURACIL AND LEUCOVORIN, FOR THE FIRST-LINE TREATMENT OF ADULT PATIENTS WITH METASTATIC PANCREATIC ADENOCARCINOMA
- I-933 REVISIONS TO THE LABELING TO EXPAND THE USE IN CERTAIN SOFT TISSUE AND CERTAIN ORTHOPEDIC SURGICAL PROCEDURES AS WELL AS REVISING THE LIMITATIONS OF USE

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

- I-934 TREATMENT OF ADULT PATIENTS WITH NEWLY DIAGNOSED PHILADELPHIA CHROMOSOME-POSITIVE ACUTE LYMPHOBLASTIC LEUKEMIA (PH+ ALL) IN COMBINATION WITH CHEMOTHERAPY
- I-935 IN COMBINATION WITH A REDUCED CALORIE DIET AND INCREASED PHYSICAL ACTIVITY TO REDUCE THE RISK OF MAJOR ADVERSE CARDIOVASCULAR EVENTS (CARDIOVASCULAR DEATH, NON-FATAL MYOCARDIAL INFARCTION, OR NON-FATAL STROKE) IN ADULTS WITH ESTABLISHED CARDIOVASCULAR DISEASE AND EITHER OBESITY OR OVERWEIGHT
- I-936 TREATMENT OF ADULT PATIENTS WITH RELAPSED OR REFRACTORY FOLLICULAR LYMPHOMA (FL), IN COMBINATION WITH OBINUTUZUMAB, AFTER TWO OR MORE LINES OF SYSTEMIC THERAPY
- I-937 EXPANSION OF THE INDICATION TO INCLUDE PRE AND PERIMENOPAUSAL WOMEN
- I-938 TREATMENT OF CHOLESTATIC PRURITUS IN PATIENTS FIVE YEARS OF AGE AND OLDER WITH PROGRESSIVE FAMILIAL INTRAHEPATIC CHOLESTASIS (PFIC)
- I-939 INDICATED FOR ACUTE TREATMENT OF MANIC OR MIXED EPISODES ASSOCIATED WITH BIPOLAR I DISORDER IN ADULTS
- I-940 TREATMENT OF CHRONIC RHINOSINUSITIS WITHOUT NASAL POLYPS (CRSSNP) IN ADULTS
- I-941 IN COMBINATION WITH PEMETREXED AND PLATINUM-BASED CHEMOTHERAPY FOR THE FIRST-LINE TREATMENT OF ADULT PATIENTS WITH LOCALLY ADVANCED OR METASTATIC NON-SMALL CELL LUNG CANCER WHOSE TUMORS HAVE EGFR EXON 19 DELETIONS OR EXON 21 L858R MUTATIONS, AS DETECTED BY AN FDA-APPROVED TEST
- I-942 EXPANSION OF INDICATION TO USE OF BICTEGRAVIR/EMTRICITABINE/TENOFOVIR ALAFENAMIDE FOR THE TREATMENT OF HIV-1 INFECTION IN VIROLOGICALLY SUPPRESSED ADULTS AND PEDIATRIC PATIENTS ON A STABLE HIV-1 TREATMENT REGIMEN WITH NO KNOWN RESISTANCE TO BICTEGRAVIR OR TENOFOVIR
- I-943 TO REDUCE THE RISK OF MYOCARDIAL INFARCTION AND CORONARY REVASCULARIZATION IN ADULTS WHO ARE UNABLE TO TAKE RECOMMENDED STATIN THERAPY (INCLUDING THOSE NOT TAKING A STATIN) WITH ESTABLISHED CARDIOVASCULAR DISEASE (CVD), OR AT HIGH RISK FOR A CVD EVENT BUT WITHOUT ESTABLISHED CVD
- I-944 EXPANDED INDICATION FOR USE AS AN ADJUNCT TO DIET, IN COMBINATION WITH OTHER LOW-DENSITY LIPOPROTEIN CHOLESTEROL (LDL-C) LOWERING THERAPIES, OR ALONE WHEN CONCOMITANT LDL-C LOWERING THERAPY IS NOT POSSIBLE, TO REDUCE LDL-C IN ADULTS WITH PRIMARY HYPERLIPIDEMIA, INCLUDING HETEROZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA (HEFH)
- I-945 EXPANDED INDICATION FOR USE AS AN ADJUNCT TO DIET, ALONE OR IN COMBINATION WITH OTHER LDL-C LOWERING THERAPIES, TO REDUCE LDL-C IN ADULTS WITH PRIMARY HYPERLIPIDEMIA, INCLUDING HETEROZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA (HEFH)
- I-946 TREATMENT OF PATIENTS 2 YEARS OF AGE AND OLDER WITH ACTIVE POLYARTICULAR JUVENILE IDIOPATHIC ARTHRITIS (PJIA) WHO HAVE HAD AN INADEQUATE RESPONSE OR INTOLERANCE TO ONE OF MORE TUMOR NECROSIS FACTOR (TNF) BLOCKERS
- I-947 FOR ADJUVANT TREATMENT IN ADULT PATIENTS FOLLOWING TUMOR RESECTION OF ANAPLASTIC LYMPHOMA KINASE (ALK)-POSITIVE NON-SMALL CELL LUNG CANCER (NSCLC) (TUMORS \geq 4 CM OR NODE POSITIVE), AS DETECTED BY AN FDA-APPROVED TEST
- I-948 USE OF VONOPRAZAN FOR THE RELIEF OF HEARTBURN ASSOCIATED WITH NON-EROSIVE GASTROESOPHAGEAL REFLUX DISEASE IN ADULTS
- I-949 FOR THE REDUCTION OF PROTEINURIA IN ADULTS WITH PRIMARY IMMUNOGLOBULIN A NEPHROPATHY (IGAN) AT RISK OF RAPID DISEASE PROGRESSION, GENERALLY A URINE PROTEIN-TO-CREATININE RATIO (UPCR) \geq 1.5 G/G
- I-950 IN COMBINATION WITH AN AROMATASE INHIBITOR FOR THE ADJUVANT TREATMENT OF ADULTS WITH HORMONE RECEPTOR (HR)-POSITIVE, HUMAN EPIDERMAL GROWTH FACTOR RECEPTOR 2 (HER2)-NEGATIVE STAGE II AND III EARLY BREAST CANCER AT HIGH RISK OF RECURRENCE
- I-951 FOR THE ADJUVANT TREATMENT OF ADULTS WITH HORMONE RECEPTOR (HR)-POSITIVE, HUMAN EPIDERMAL GROWTH FACTOR RECEPTOR 2 (HER2)-NEGATIVE STAGE II AND III EARLY BREAST CANCER AT HIGH RISK OF RECURRENCE
- I-952 TREATMENT OF ADULT PATIENTS WITH LOCALLY ADVANCED, UNRESECTABLE (STAGE III) NON-SMALL CELL LUNG CANCER (NSCLC) WHOSE DISEASE HAS NOT PROGRESSED DURING OR FOLLOWING CONCURRENT OR SEQUENTIAL PLATINUM-BASED CHEMORADIATION THERAPY AND WHOSE TUMORS HAVE EGFR EXON 19 DELETIONS OR EXON 21 L858R MUTATIONS, AS DETECTED BY AN FDA-APPROVED TEST
- I-953 TREATMENT OF ADULT PATIENTS WITH NEWLY DIAGNOSED PHILADELPHIA CHROMOSOME-POSITIVE CHRONIC MYELOID LEUKEMIA (PH+ CML) IN CHRONIC PHASE (CP)

PATENT AND EXCLUSIVITY TERMS

ADB 41 of 225

EXCLUSIVITY INDICATION

- I-954 TREATMENT OF ADULT PATIENTS WITH PREVIOUSLY TREATED PHILADELPHIA CHROMOSOME-POSITIVE CHRONIC MYELOID LEUKEMIA (PH+ CML) IN CHRONIC PHASE (CP)
- I-955 TREATMENT OF OVERACTIVE BLADDER (OAB) WITH SYMPTOMS OF URGE URINARY INCONTINENCE, URGENCY, AND URINARY FREQUENCY IN ADULT MALES ON PHARMACOLOGICAL THERAPY FOR BENIGN PROSTATIC HYPERPLASIA (BPH)
- I-956 TREATMENT OF ATOPIC DERMATITIS IN ADULT AND PEDIATRIC PATIENTS 2 YEARS OF AGE AND OLDER

EXCLUSIVITY MISCELLANEOUS

- M-1 INFORMATION REGARDING SUPERIORITY CLAIM OVER RANITIDINE FOR DAY AND NIGHT HEARTBURN ADDED TO CLINICAL STUDIES SECTION
- M-2 APPROVAL FOR ADDITION TO CLINICAL PHARMACOLOGY SECTION OF THE LABEL REGARDING (1) IMPROVEMENT IN BONE MINERAL DENSITY IN CHILDHOOD-ONSET ADULT GROWTH HORMONE DEFICIENT PATIENTS AND (2) INCREASES IN SERUM ALKALINE PHOSPHATASE
- M-3 ADDITION OF EFFICACY AND SAFETY INFORMATION IN WHICH FOSAMAX WAS USED CONCOMITANTLY WITH ESTROGEN ALONE OR WITH ESTROGEN PLUS PROGESTIN
- M-4 CHANGES TO PEDIATRIC USE SECTION TO PROVIDE INFORMATION REGARDING SAFETY AND EFFICACY IN PEDIATRIC PATIENTS AS YOUNG AS 2 YEARS OLD
- M-5 INFORMATION REGARDING EFFECTS IN PATIENTS WITH ASTHMA ON CONCOMITANT INHALED CORTICOSTEROIDS IN CLINICAL PHARMACOLOGY SECTION
- M-6 ADDITIONAL INFORMATION REGARDING CLINICAL STUDIES DONE WITH GLUCOPHAGE/GLYBURIDE COMBINATION ADDED TO CLINICAL PHARMACOLOGY AND DOSING AND ADMINISTRATION
- M-7 CLINICAL PHARMACOLOGY IN PEDIATRIC PATIENTS; DOSAGE AND ADMINISTRATION INFORMATION
- M-8 ADDITIONAL INFORMATION FOR THE USE OF SONATA CAPSULES FOR UP TO 5 WEEKS (35 NIGHTS) OF TREATMENT IN A CONTROLLED TRIAL SETTING
- M-9 ADDITION TO THE CLINICAL STUDIES SECTION OF THE LABELING OF TEXT AND TWO TABLES CONTAINING INFORMATION FOR THE PRESCRIBING PHYSICIAN ON BLOOD PRESSURE, HEART RATE, AND HEART RATE VARIABILITY
- M-10 INFORMATION REGARDING MAINTENANCE OF AN ANTIDEPRESSANT EFFECT UP TO 1 YEAR OF DOSING
- M-11 USE FOR LONG-TERM TREATMENT OF POSTTRAUMATIC STRESS DISORDER
- M-12 NEW LANGUAGE FOR PEDIATRIC USE
- M-13 INFORMATION FROM PEDIATRIC STUDIES ADDED TO CLINICAL PHARMACOLOGY, PRECAUTIONS, AND DOSAGE AND ADMINISTRATION
- M-14 ADDITIONAL CLINICAL TRIAL INFORMATION ADDED TO PEDIATRIC USE SUBSECTION
- M-15 LONGER TERM EFFICACY INFORMATION FOR RISPERIDONE IN THE TREATMENT OF SCHIZOPHRENIA
- M-16 CHANGE IN WORDING OF THE PEDIATRIC SECTION OF THE PACKAGE INSERT
- M-17 INFORMATION REGARDING USE OF ULTANE IN PEDIATRIC PATIENTS WITH CONGENITAL HEART DISEASE
- M-18 INFORMATION DENOTING THE EFFICACY OF REMERON IN MAINTAINING A RESPONSE IN PATIENTS WITH MAJOR DEPRESSIVE DISORDER (MDD)
- M-19 INFORMATION REGARDING USE IN PEDIATRIC PATIENTS TWO YEARS OF AGE AND OLDER
- M-20 LABELING REVISIONS RELATED TO MCCUNE ALBRIGHT SYNDROME
- M-21 COMPARISON DATA ON THE ANTIHYPERTENSIVE EFFECTS OF ATACAND AND COZAAR
- M-22 CHANGE IN TIME TO ONSET OF ACTION
- M-23 INFORMATION REGARDING ELIMINATION ADDED TO CLINICAL PHARMACOLOGY, STUDY RESULTS IN PATIENTS WITH HEPATIC AND RENAL IMPAIRMENT
- M-24 INFORMATION ON RESULTS OF A LONG TERM LONGITUDINAL GROWTH STUDY AND PEDIATRIC SAFETY INFORMATION
- M-25 ADDITIONAL SAFETY AND PHARMACOKINETICS INFORMATION IN CHILDREN 6 MONTHS TO LESS THAN 6 YEARS OF AGE ADDED TO PACKAGE INSERT

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY MISCELLANEOUS**

- M-26 INCORPORATION OF INFORMATION CONTAINED IN THE PEG-INTRON PACKAGE INSERT INTO THE REBETOL PACKAGE INSERT AND MEDGUIDE-PEG-INTRON WAS APPROVED FOR USE IN COMBINATION WITH REBETOL FOR TREATMENT OF CHRONIC HEPATITIS C VIRUS INFECTION ON 8/7/01
- M-27 INFORMATION DESCRIBING ASPIRIN ENDOSCOPY STUDY AND THE MAXIMUM RECOMMENDED DOSE FOR PATIENTS WITH MODERATE HEPATIC INSUFFICIENCY
- M-28 INFORMATION FROM A STUDY IN PEDIATRIC PATIENTS IN ASSOCIATION WITH A NEUROLOGICAL CONDITION
- M-29 LABELING CHANGES TO PROVIDE INFORMATION IN THE MANAGEMENT OF OBESITY IN ADOLESCENTS AGED 12 TO 16 YEARS
- M-30 CHANGES TO CLINICAL PHARMACOLOGY, PRECAUTIONS, AND DOSAGE AND ADMINISTRATION SECTIONS OF LABELING CONCERNING USE OF LOTENSIN IN PEDIATRIC PATIENTS WITH HYPERTENSION
- M-31 INFORMATION FOR USE IN PEDIATRIC PATIENTS WITH CHRONIC KIDNEY DISEASE STAGE 5 (END-STAGE RENAL DISEASE)
- M-32 ADDITIONAL LANGUAGE TO CLINICAL PHARMACOLOGY AND CLINICAL STUDIES
- M-33 INFORMATION FOR USE OF ADVAIR DISKUS 100/50 IN CHILDREN 4 TO 11 YEARS OF AGE WITH ASTHMA
- M-34 EXPANDED INFORMATION TO PEDIATRIC USE SUBSECTION OF LABELING IN RESPONSE TO PEDIATRIC WRITTEN REQUEST
- M-35 ADDITIONAL INFORMATION REGARDING CLINICAL STUDIES DONE WITH ACTOS IN COMBINATION WITH METFORMIN, A SULFONYLUREA, OR INSULIN ADDED TO CLINICAL PHARMACOLOGY
- M-36 ADDITION OF INFORMATION TO CLINICAL STUDIES REGARDING PREVENTION OF CARDIOVASCULAR DISEASE
- M-37 INFORMATION ADDED TO THE LABELING THAT DETAILS INFORMATION RELATIVE TO STUDIES DONE IN PEDIATRIC POPULATIONS IN THE CLINICAL PHARMACOLOGY AND PEDIATRIC USE SUBSECTIONS
- M-38 SAFETY AND IOP-LOWERING EFFECTS OF TRUSOPT HAVE BEEN DEMONSTRATED IN PEDIATRIC PATIENTS IN A 3 MONTH, MULTI-CENTER DOUBLE MASKED ACTIVE-TREATMENT-CONTROLLED TRIAL
- M-39 FOR LABELING CHANGES BASED ON RESULTS OF THE SPD422-202 CLINICAL STUDY REPORT (CSR) SUBMITTED IN RESPONSE TO THE WRITTEN REQUEST
- M-40 ADDITIONAL INFORMATION REGARDING CLINICAL STUDIES PERFORMED IN PEDIATRIC PATIENTS WITH LEUKEMIA ADDED TO PRECAUTIONS
- M-41 REVISION TO THE PEDIATRIC USE PRECAUTIONS OF THE PRESCRIBING INFORMATION TO INCORPORATE THE RESULTS FROM THE CAPPS-169 STUDY ENTITLED "THE EFFECT OF ORTHO TRICYCLEN ON BONE MINERAL DENSITY IN PEDIATRIC SUBJECTS WITH ANOREXIA NERVOSA"
- M-42 ADDITION OF A GERIATRIC USE SUBSECTION TO THE PRECAUTIONS SECTION OF THE PACKAGE INSERT AND GERIATRIC DOSING INFORMATION
- M-43 INCLUSION OF RESULTS OF STUDY "PLACEBO-CONTROLLED STUDY TO EVALUATE SAFETY AND PILOT EFFICACY OF ILOPROST AS ADD ON THERAPY WITH BOSENTAN IN SUBJECTS WITH PULMONARY ARTERIAL HYPERTENSION"
- M-44 CLINICAL INFORMATION ADDED TO THE PEDIATRIC USE SUBSECTION OF PRECAUTIONS REGARDING THE USE OF NOVOLOG IN ADOLESCENTS WITH TYPE I DIABETES AGE 6 TO 18
- M-45 INFORMATION ADDED TO CLINICAL TRIALS SECTION OF LABELING, "EFFECTS OF HUMATROPE TREATMENT IN ADULTS WITH GROWTH HORMONE DEFICIENCY"
- M-46 PROVISION OF RESULTS OF STUDY AND PROPOSED REVISIONS TO PACKAGE INSERT SEE SECTION ON CARDIAC ELECTROPHYSIOLOGY
- M-47 PROVIDES FOR USE OF ANTARA WITHOUT REGARD TO MEALS
- M-48 CHANGES TO THE LABELING DESCRIBING THE RESULTS OF A STUDY OF THE USE OF NOVOLOG MIX 70/30 WITH ORAL ANTIDIABETIC AGENTS IN PATIENTS WITH TYPE 2 DIABETES
- M-49 CLINICAL DATA ADDED TO THE CLINICAL PHARMACOLOGY SECTION REGARDING EFFECT OF SINGULAIR ON GROWTH RATES IN PREPUBERTAL CHILDREN
- M-50 NEW INFO TO THE CLINICAL STUDIES, ADULT GROWTH HORMONE DEFICIENCY (GHD) SUBSECTION OF THE NUTROPIN AQ PACKAGE INSERT DESCRIBING THE EFFECTS OF SOMATROPIN ON VISCERAL ADIPOSE TISSUE IN THE ADULT GROWTH HORMONE DEFICIENT PATIENT POPULATION
- M-51 INFORMATION ADDED TO LABELING REGARDING OSTEOGENESIS IMPERFECTA STUDY

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY MISCELLANEOUS**

- M-52 INFORMATION ADDED TO THE CLINICAL PHARMACOLOGY/CLINICAL STUDIES SECTION REGARDING THE USE OF RISEDRONATE ADMINISTERED ONCE A WEEK IN THE PREVENTION OF OSTEOPOROSIS IN POSTMENOPAUSAL WOMEN
- M-53 FOR LABELING CHANGES TO THE QUALITY OF LIFE (QOL) STATEMENT IN THE APPROVED PACKAGE INSERT
- M-54 INFORMATION FROM PEDIATRIC STUDIES ADDED TO LABEL
- M-55 INFORMATION ON RESULTS OF A STUDY OF THE USE OF SANDOSTATIN LAR DEPOT IN PEDIATRIC PATIENTS WITH HYPOTHALAMIC OBESITY.
- M-56 INFORMATION ADDED TO CLINICAL TRIAL SECTION WITH INFORMATION ON "GEMINI" TRIAL
- M-57 CLINICAL DATA ADDED TO THE CLINICAL PHARMACOLOGY SECTION REGARDING THE PHARMACOKINETICS OF EZETIMIBE IN ASIAN SUBJECTS
- M-58 CHANGES TO THE CLINICAL STUDIES, PRIMARY HYPERCHOLESTEROLEMIA, VYTORIN SUBSECTION OF THE PACKAGE INSERT TO ADD EFFICACY DATA FOR THE EZETIMIBE/SIMVASTATIN COMBINATION PRODUCT AND FOR AN ATORVASTATIN PRODUCT ON LDL-C AND OTHER LIPID PRMTRS
- M-59 RESULTS OF THE T20-310 STUDY WHICH EVALUATED THE PHARMACOKINETICS, SAFETY, AND ANTIVIRAL ACTIVITY OF FUZEON IN TREATMENT EXPERIENCED PEDIATRIC SUBJECTS AND ADOLESCENTS WAS ADDED TO THE PEDIATRIC SUBSECTION OF PRECAUTIONS
- M-60 CHANGES TO CLINICAL STUDIES, PRIMARY HYPERCHOLESTEROLEMIA, TO ADD EFFICACY DATA FOR THE EZETIMIBE/SIMVASTATIN COMBINATION PRODUCT AND FOR A ROSUVASTATIN PRODUCT ON LDL-C AND OTHER LIPID PARAMETERS IN PATIENTS WTH HYPERCHOLESTEROLEMIA
- M-61 REVISIONS TO LABELING BASED ON DATA SUBMITTED IN RESPONSE TO PEDIATRIC WRITTEN REQUEST
- M-62 CLINICAL INFORMATION FROM ONE CLINICAL STUDY INVESTIGATING THE USE OF AVANDAMET PLUS INSULIN IN PATIENTS WITH TYPE 2 DIABETES MELLITUS WHO HAVE NOT ACHIEVED ADEQUATE GLYCEMIC CONTROL WITH PREVIOUS ANTI-DIABETIC THERAPIES
- M-63 DETAILED INFORMATION ON AN INCONCLUSIVE PEDIATRIC STUDY
- M-64 CHANGES TO CLINICAL PHARMACOLOGY DETAILING STUDY RESULTS
- M-65 ADDITION OF INFORMATION TO LABEL TO INCLUDE INFORMATION REGARDING USE IN PATIENTS WITH HIV-ASSOCIATED ADIPOSE REDISTRIBUTION SYNDROME (HARS)
- M-66 USE IN SPECIFIC POPULATIONS - PATIENTS WITH CONCOMITANT ILLNESS SUBSECTION OF THE LABELING REGARDING USE OF STRATTERA IN PATIENTS WITH ADHD WHO HAVE COMORBID TIC DISORDER
- M-67 INDICATION EXPANDED TO INCLUDE PATIENTS ON PERITONEAL DIALYSIS
- M-68 DESCRIPTION OF RESULTS OF STUDY OF INITIAL THERAPY IN COMBINATION WITH METFORMIN WHEN DIET AND EXERCISE DO NOT PROVIDE GLYCEMIC CONTROL
- M-69 RESULTS OF STUDY OF COMBINATION THERAPY AND NON-INFERIORITY STUDY
- M-70 PROVISION OF INFORMATION OF THE RESULTS OF A PHASE 2 RANDOMIZED TRIAL OF SPRYCEL 70MG TWICE DAILY OR IMATINIB 800MG DAILY
- M-71 REVISIONS TO PROVIDE FOR RESULTS OF MAINTENANCE DATA IN ADULT PATIENTS WITH MAJOR DEPRESSIVE DISORDER
- M-72 INFORMATION ABOUT USE OF INSPIRA (EPLERENONE) FOR HYPERTENSION IN PEDIATRIC PATIENTS
- M-73 NEW INFORMATION ADDED REGARDING THE TUMOR SHRINKING POTENTIAL OF SANDOSTATIN LAR DEPOT INJECTION ON GH - SECRETING PITUITARY ADENOMAS
- M-74 REVISIONS TO CLINICAL STUDIES - CHILDREN AND ADOLESCENTS BASED ON CLINICAL TRIAL DATA TO SUPPORT A DURATION OF ACTION CLAIM UP TO 12 HOURS
- M-75 PROVISION FOR USE OF ARGATROBAN IN CERTAIN PEDIATRIC PATIENTS WITH HEPARIN-INDUCED THROMBOCYTOPENIA (HIT) OR HEPARIN-INDUCED THROMBOCYTOPENIA WITH THROMBOSIS (HITTS)
- M-76 REMOVAL OF SCREEN REQUIREMENT IN PTS WITH G6PD DEFICIENCY PRIOR TO INITIATING ACZONE TREATMENT; REMOVAL OF BLOOD COUNT & RETICULOCYTE MONITORING DURING TREATMENT IN G6PD DEFICIENT PTS AND IN PATIENTS WITH HISTORY OF ANEMIA
- M-77 USE IN COMBINATION WITH THE NEW AKTILITE CL128 LAMP FOR THE TREATMENT OF THIN AND MODERATELY THICK, NON-HYPERKERATOTIC, NON-PIGMENTED ACTINIC KERATOSES OF THE FACE AND SCALP IN IMMUNOCOMPETENT PATIENTS
- M-78 CLINICAL TRIAL INFO ON USE OF STRATTERA IN PATIENTS WITH ATTENTION DEFICIT HYPERACTIVITY DISORDER (ADHD) AND COMORBID ANXIETY DISORDER WITHOUT CAUSING

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY MISCELLANEOUS**

WORSENING OF ANXIETY

- M-79 LABELING REVISIONS RELATED TO SMOKING AND ERLOTINIB EXPOSURE
- M-80 ADDITIONAL TIME POINT OF 30 MINUTES (0.5 HOUR) IN CHILDREN AGED 6-12 YEARS WITH A DIAGNOSIS OF ATTENTION-DEFICIT HYPERACTIVITY DISORDER (ADHD)
- M-81 ADDITIONAL INFO FOR PEDIATRIC USE FOR CASODEX (STUDIED IN COMBINATION WITH ARIMIDEX) IN THE PEDIATRIC POPULATION, SPECIFICALLY BOYS WITH FAMILIAL MALE-LIMITED PRECOCIOUS PUBERTY (TESTOTOXICOSIS)
- M-82 LABELING REVISIONS RELATED TO CLINICAL STUDIES
- M-83 ADDITIONAL INFORMATION ADDED TO LABELING REGARDING ESTABLISHMENT OF EFFICACY IN ADDITIONAL CLINICAL TRIALS AND ONE MAINTENANCE TRIAL
- M-84 STUDY INFORMATION ADDED TO LABEL REGARDING BONE MINERAL DENSITY
- M-85 INFORMATION ADDED TO LABELING REGARDING USE OF PREVACID IN PATIENTS LESS THAN 1 YEAR WITH SYMPTOMATIC GERD
- M-86 LABELING CHANGES SUBMITTED IN RESPONSE TO PEDIATRIC WRITTEN REQUEST FOR INFANTS AGES BIRTH TO 11 MONTH INCLUSIVE REFLECTING LACK OF EFFICACY FOR GERD INDICATION FOR THIS PATIENT POPULATION
- M-87 INCLUSION OF RESULTS FROM TWO DRUG INTERACTION STUDIES WITH LIPITOR AND CRESTOR IN CLINICAL PHARMACOLOGY SECTION
- M-88 ADDITION OF INFORMATION REGARDING ABUSE POTENTIAL OF CONCERTA VERSUS IMMEDIATE-RELEASE METHYLPHENIDATE
- M-89 PROVIDES FOR REVISIONS TO MULTIPLE SECTIONS OF THE PACKAGE INSERT TO REFLECT RESULTS OF CLINICAL TRIALS 205.235 (UPLIFT) AND 205.266 (VA STUDY) IN SUPPORT OF EXACERBATION CLAIM
- M-90 LABELING CHANGES BASED ON DATA FROM CLINICAL STUDIES NV20235 AND NV20236 STUDIES OF SEASONAL PROPHYLAXIS OF INFLUENZA IN IMMUNOCOMPROMISED PATIENTS AND CHILDREN AGES 1-12
- M-91 UPDATED LABELING BASED UPON STUDY: A SINGLE-DOSE, SINGLE-BLIND, PLACEBO-AND MOXIFLOXACIN-CONTROLLED 2-PERIOD, RANDOMIZED, CROSSOVER, 3RD PERIOD SEQUENTIAL STUDY OF SIDE EFFECTS OF TEMSIROLIMUS ON CARDIAC REPOLARIZATION IN HEALTHY SUBJECTS
- M-92 UPDATES TO THE PACKAGE INSERT BASED UPON THE TRIAL ENTITLED "A PHASE I PHARMACOKINETIC AND PHARMACODYNAMIC STUDY OF TEMSIROLIMUS IN PATIENTS WITH ADVANCED MALIGNANCIES AND NORMAL AND IMPAIRED LIVER FUNCTION"
- M-93 EXPANSION OF LABELING TO INCLUDE INFORMATION ON SAFETY AND EFFICACY OF CREON IN PATIENTS AGES 7 YEARS THROUGH 11 YEARS WITH PANCREATIC EXOCRINE INSUFFICIENCY DUE TO CYSTIC FIBROSIS
- M-94 INFO ADDED TO LABEL RELATED TO NEWLY DIAGNOSED PHILADELPHIA CHROMOSOME POSITIVE (PH+) CHRONIC MYELOID LEUKEMIA IC CHRONIC PHASE
- M-95 INFORMATION FOR TREATMENT OF CHRONIC HEPATITIS B (CHB) IN ADULT PATIENTS WITH DECOMPENSATED LIVER DISEASE BASED ON DATA FROM CLINICAL TRIAL GS-US-174-0108
- M-96 UPDATED INFORMATION IN THE CLINICAL STUDIES SECTION RELATED TO THE LOSS AND RECOVERY OF BONE MINERAL DENSITY IN ADOLESCENT GIRLS DURING AND FOLLOWING THE USE OF DEPO-PROVERA CONTRACEPTIVE INJECTION
- M-97 LABELING CHANGES IN RESPONSE TO PEDIATRIC STUDIES - NOT INDICATED FOR USE IN PEDIATRIC POPULATION
- M-98 NEW INFORMATION FROM A STUDY WHICH EVALUATED THE SAFETY AND EFFICACY OF FAMVIR IN TREATING RECURRENT GENITAL HERPES IN IMMUNOCOMPETENT BLACK/AFRICAN AMERICAN SUBJECTS.
- M-99 ADDITION OF FINDINGS FROM A SINGLE PEDIATRIC CLINICAL TRIAL (P04292) OF NASONEX NASAL SPRAY IN THE TREATMENT OF NASAL POLYPS IN PATIENTS 6 TO <18 YEARS OF AGE TO THE PACKAGE INSERT.
- M-100 INFORMATION ADDED TO LABEL BASED UPON COMPLETED CLINICAL TRIAL REPORTS
- M-101 INCLUSION OF DATA FROM AN ADDITIONAL 19 SUBJECTS WITH HYPERCALCEMIA FROM PARATHYROID CARCINOMA TO THE INFORMATION CURRENTLY PRESENTED IN THE LABEL
- M-102 INFORMATION FROM PEDIATRIC STUDY REPORT ML16633, "INTRAVENOUS GRANISETRON (KYTRIL) IN THE PREVENTION OF POST-OPERATIVE NAUSEA AND VOMITING (PONV) IN PEDIATRIC SUBJECTS UNDERGOING TONSILLECTOMY OR ADENOTONSILLECTOMY."
- M-103 SAFETY, EFFICACY AND PHARMACOKINETIC INFO FOR FASLODEX IN THE PEDIATRIC

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY MISCELLANEOUS**

- POPULATION, SPECIFICALLY FOR GIRLS WITH PROGRESSIVE PRECOCIOUS PUBERTY ASSOCIATED WITH MCCUNE-ALBRIGHT SYNDROME ADDED TO THE PEDIATRIC USE SECTION OF THE LABELING
- M-104 INFORMATION ADDED TO DOSING AND ADMINISTRATION REGARDING A 26 WEEK STUDY
- M-105 NEW LANGUAGE ADDED TO CLINICAL STUDIES REGARDING USE IN SMOKERS WITH CARDIOVASCULAR DISEASE, CHRONIC OBSTRUCTIVE PULMONARY DISEASE, AND USE ACCORDING TO AN ALTERNATIVE SET OF DIRECTIONS FOR SETTING A QUIT DATE
- M-106 ADDITION OF THE T1-WEIGHTED GD-ENHANCED LESION EFFICACY VARIABLE IN THE CLINICAL STUDIES SECTION 14 OF THE PACKAGE INSERT
- M-107 INFORMATION TO THE CLINICAL STUDIES SECTION OF THE LUPRON DEPOT-PED 1-MONTH BASED UPON THE PHASE 3/4 COMPLETED CLINICAL STUDY REPORT FOR STUDY M90-516 ENTITLED "STUDY OF LUPRON DEPOT IN THE TREATMENT OF CENTRAL PRECOCIOUS PUBERTY".
- M-108 CHANGES ARE BASED ON RESULTS FROM STUDY CV181057
- M-110 CHANGES TO THE PACKAGE INSERT TO REFLECT THE RESULTS OF THE STUDY OF HEART AND RENAL PROTECTION (SHARP) TRIAL
- M-111 LABELING CHANGES BASED ON STUDY HW80-EW-GWCI ENTITLED A PLACEBO AND POSITIVE CONTROLLED STUDY OF THE ELECTROPHYSIOLOGICAL EFFECTS OF A SINGLE 10 MCG DOSE OF EXENATIDE ON THE 12 LEAD ELECTROCARDIOGRAM QT INTERVAL IN HEALTHY SUBJECTS
- M-112 REVISIONS TO THE PEDIATRIC USE SECTION OF THE PACKAGE INSERT TO ADD INFORMATION FROM A PEDIATRIC STUDY IN PATIENTS AGED 12 YEARS TO LESS THAN 18 YEARS OF AGE WITH RECURRENT HERPES LABIALIS
- M-113 LABELING CHANGES BASED ON STUDY H80-US-GWCO ENTITLED A RANDOMIZED TRIAL COMPARING EXENATIDE WITH PLACEBO IN SUBJECTS WITH TYPE 2 DIABETES ON INSULIN GLARGINE WITH OR WITHOUT ORAL ANTIHYPERGLYCEMIC MEDICATIONS
- M-114 CHANGES IN SECTION 14 OF THE PACKAGE INSERT TO INCLUDE DATA FROM THE SWITCHMRK STUDIES (SWITCH OF SUPPRESSED SUBJECTS FROM LOPINAVIR/RITONAVIR TO RALTEGRAVIR)
- M-115 REVISIONS TO THE PI BASED ON RESULTS FROM STUDY NN2211-1842, ENTITLED THE EFFECT OF INSULIN DETEMIR IN COMBINATION WITH LIRAGLUTIDE AND METFORMIN COMPARED TO LIRAGLUTIDE AND METFORMIN IN SUBJECTS WITH TYPE 2 DIABETES
- M-116 LABELING CHANGES BASED ON RESULTS FROM CLINICAL STUDY 01-06-TL-OPIMET-008
- M-117 ADDITION OF RESULTS OF PEDIATRIC TRIAL TO LABEL
- M-118 LABELING CHANGES BASED UPON SAFETY AND EFFICACY RESULTS FROM TRIAL 1218.36
- M-119 LABELING CHANGES REGARDING MISSED DOSES
- M-120 CHANGES TO CLINICAL TRIALS DETAILING STUDY RESULTS
- M-121 LABELING CHANGES BASED UPON SAFETY AND EFFICACY RESULTS FROM TRIAL 1218.43
- M-122 LABELING CHANGES TO INCLUDE THE RESULTS OF THE PARAMOUNT TRIAL
- M-123 UPDATED RESULTS OF OVERALL SURVIVAL FROM 'CONFIRM' STUDY
- M-124 LONG TERM SAFETY AND EFFICACY DATA FROM STUDY CLDT600A2303 FOR SUBJECTS PREVIOUSLY ENROLLED IN THE ORIGINAL TWO YEAR GLOBE (NV-02B-007/CLDT600A2302) AND NV02B-015 STUDIES WHO CONTINUED TELBIVUDINE TREATMENT FOR UP TO 208 WEEKS
- M-125 LABELING CHANGES TO INCLUDE LACK OF EFFICACY IN CHILDREN 6 MONTHS TO 4 YEARS OF AGE
- M-126 UPDATES TO THE CLINICAL STUDIES SECTION 14, OF THE PACKAGE INSERT (PI), WITH THE RESULTS OF CLINICAL TRIAL P06086
- M-127 REVISIONS TO THE PEDIATRIC USE SECTION OF THE PACKAGE INSERT TO REFLECT THE RESULTS FROM CLINICAL STUDY C-10-004
- M-128 CLINICAL TRIAL STUDY RESULTS
- M-129 RESULTS OF A CLINICAL STUDY REPORT WHICH ASSESSES THE SAFETY AND EFFICACY IN CHILDREN AGES 6 TO 12 YEARS OF AGE
- M-130 ADDITION OF INFORMATION ON LONG-TERM TREATMENT WITH VPRIV IN THE CLINICAL TRIALS SECTION OF THE PACKAGE INSERT
- M-131 INFORMATION FROM STUDIES CONDUCTED IN PEDIATRIC PATIENTS WITH NEWLY DIAGNOSED NON-DISSEMINATED DIFFUSED INTRINSIC BRAINSTEM GLIOMAS
- M-132 REVISIONS TO THE CLINICAL TRIALS SECTION IN THE INOMAX LABEL TO REFLECT RESULTS FROM THE PEDIATRIC STUDY REPORTS
- M-133 INFORMATION ADDED TO THE LABELING REGARDING THE ADDITION OF SILDENAFIL TO

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY MISCELLANEOUS**

BOSENTAN THERAPY

- M-134 ADDITIONAL INFORMATION REGARDING CLINICAL STUDIES PERFORMED WITH SAXAGLIPTIN IN COMBINATION WITH METFORMIN AND A SULFONYLUREA ADDED TO THE LABELING
- M-135 ADDITION OF INFORMATION TO THE CLINICAL STUDIES - RADIOGRAPHIC RESPONSE SECTION OF THE PACKAGE INSERT
- M-136 ADDITIONAL INFORMATION ADDED TO THE USE IN SPECIFIC POPULATIONS SECTION OF THE LABELING REGARDING POST-OPERATIVE NAUSEA AND VOMITING STUDIES IN PEDIATRIC PATIENTS
- M-137 LABELING REVISIONS RESULTING FROM A MAINTENANCE TRIAL IN PEDIATRIC PATIENTS WITH IRRITABILITY ASSOCIATED WITH AUTISTIC DISORDER
- M-138 INFORMATION ADDED TO THE 8.4 PEDIATRIC USE SECTION ON THE USE OF MEMANTINE IN CHILDREN AGES 6-12 YEARS WITH AUTISM SPECTRUM DISORDER
- M-139 INFORMATION ADDED TO THE DOSING AND ADMINISTRATION SECTION OF THE PACKAGE INSERT REGARDING RETREATMENT WITH VELCADE FOR PATIENTS WITH MULTIPLE MYELOMA
- M-140 INFORMATION ADDED TO THE PEDIATRIC USE SECTION OF THE LABELING REGARDING USE OF LATISSE IN PATIENTS WHO WERE POST-CHEMOTHERAPY OR HAD ALOPECIA AREATA, AND ADOLESCENTS WHO HAD HYPERTRICHOSIS WITH NO ASSOCIATED MEDICAL CONDITION
- M-141 REVISIONS TO THE PEDIATRIC USE SECTION OF THE LABELING TO INCORPORATE STUDY RESULTS FOR TREATMENT OF MAJOR DEPRESSIVE DISORDER IN ADOLESCENTS (AGES 12-17)
- M-142 ADDITIONS TO THE LABELING DESCRIBING RESULTS FROM STUDY H6P-MC-HDAY
- M-143 INFORMATION ADDED TO THE LABELING REGARDING THE SAFETY AND EFFICACY OF VARENICLINE FOR SMOKING CESSATION IN PATIENTS WITH CURRENT OR PAST HISTORY OF MAJOR DEPRESSIVE DISORDER
- M-144 INFORMATION ADDED TO THE LABELING REGARDING THE SAFETY AND EFFICACY OF VARENICLINE FOR SMOKING CESSATION IN PATIENTS WHO HAD BEEN PREVIOUSLY TREATED WITH VARENICLINE
- M-145 ADDITION OF INFORMATION ABOUT LONG-TERM TREATMENT OF PULMONARY ARTERIAL HYPERTENSION TO THE CLINICAL STUDIES SECTION OF THE LABELING
- M-146 INFORMATION ADDED TO THE CLINICAL STUDIES SECTION ON INITIAL COMBINATION THERAPY WITH LINAGLIPTIN AND METFORMIN VS. LINAGLIPTIN MONOTHERAPY IN TREATMENT NAIVE PATIENTS
- M-147 OTC USE FOR TEMPORARY RELIEF OF OCULAR SYMPTOMS DUE TO HAY FEVER OR OTHER UPPER RESPIRATORY ALLERGIES
- M-148 LABELING CHANGES BASED ON STUDY H80-EW-GWDM
- M-149 INFORMATION ADDED TO THE LABELING REGARDING MAINTENANCE MONOTHERAPY FOR ADHD
- M-150 ADDITION OF THE RESULTS OF A CONTROLLED CLINICAL STUDY TREATING ADULT PATIENTS WITH SCHIZOPHRENIA EXPERIENCING AN ACUTE RELAPSE
- M-151 REVISIONS TO THE LABELING BASED ON THE OUTCOMES OF PEDIATRIC STUDIES CONDUCTED TO ASSESS THE SAFETY AND EFFICACY OF XOPENEX IN SUBJECTS LESS THAN 6 YEARS OF AGE
- M-152 INFORMATION ADDED TO THE CLINICAL PHARMACOLOGY SECTION OF THE LABELING REGARDING A SAFETY STUDY IN PEDIATRIC SUBJECTS AGES 6 MONTHS TO 4 YEARS OF AGE WITH AN ACTIVE HEAD LICE INFESTATION
- M-153 ADDITION OF INFORMATION REGARDING THE INTRANASAL ABUSE POTENTIAL OF OXYCONTIN
- M-154 UPDATE TO THE LABELING TO REFLECT THE RESULTS OF A LONG-TERM MAINTENANCE TREATMENT STUDY OF ADHD IN CHILDREN AND ADOLESCENTS AGES 6-17.
- M-155 ADDITION OF CLINICAL FINDINGS FROM AN OBSERVATIONAL STUDY IN A PEDIATRIC AGE GROUP GREATER THAN 2 MONTHS TO 18 YEARS IN SECTION 8.4 PEDIATRIC USE OF THE PACKAGE INSERT
- M-156 UPDATE TO THE LABELING WITH INFORMATION REGARDING A CLINICAL TRIAL IN CHILDREN LESS THAN 4 YEARS OF AGE.
- M-157 INFORMATION ADDED TO THE LABELING REGARDING THE SAFETY AND EFFICACY OF DAPAGLIFLOZIN 10MG ONCE DAILY IN PATIENTS WITH TYPE 2 DIABETES WHO HAVE INADEQUATE GLYCEMIC CONTROL ON A BACKGROUND COMBINATION OF METFORMIN AND SULFONYLUREA
- M-158 UPDATES TO THE LABELING TO REFLECT SAFETY RESULTS FROM CLINICAL TRIALS IN SCHIZOPHRENIA ADOLESCENT PATIENTS AGED 12 TO 17 YEARS

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY MISCELLANEOUS**

- M-159 ADDITION OF PED SAFETY INFORMATION DERIVED FROM A MAINTENANCE TREATMENT STUDY OF BIPOLAR 1 DISORDER TO DELAY THE TIME TO OCCURRENCE OF MOOD EPISODES IN PATIENTS (> THAN OR = TO 13 YRS OF AGE) TREATED FOR ACUTE MOOD EPISODES WITH STANDARD THERAPY
- M-160 UPDATED LABELING WITH DATA FROM A RANDOMIZED, DOUBLE-BLIND ACTIVE-CONTROLLED STUDY COMPARING EMPAGLIFLOZIN TO GLIMEPIRIDE IN PATIENTS WITH TYPE 2 DIABETES AND INSUFFICIENT GLYCEMIC CONTROL DESPITE METFORMIN TREATMENT
- M-161 UPDATED LABELING WITH DATA FROM A RANDOMIZED, DOUBLE-BLIND, PLACEBO-CONTROLLED STUDY OF EMPAGLIFLOZIN IN PATIENTS WITH TYPE 2 DIABETES MELLITUS AND INSUFFICIENT GLYCEMIC CONTROL ON A MULTIPLE DAILY INJECTION INSULIN REGIMEN ALONE OR WITH METFORMIN
- M-162 INCLUSION OF EFFICACY AND SAFETY DATA TO THE PRESCRIBING INFORMATION OF BYDUREON BASED ON STUDY GWDE
- M-163 INFORMATION ADDED TO THE LABELING REGARDING PREVIOUSLY UNTREATED ALK-POSITIVE METASTATIC NON SMALL CELL LUNG CANCER (NSCLC)
- M-164 REVISES THE CLINICAL TRIALS SECTION OF THE PRESCRIBING INFORMATION TO INCORPORATE THE RESULTS FROM STUDY E7273-G000-401 ENTITLED "PHASE IV RANDOMIZED STUDY OF TWO DOSE LEVELS OF TARGRETIN CAPSULES IN SUBJECTS WITH REFRACTORY CUTANEOUS T-CELL LYMPHOMA"
- M-165 PROVIDES FOR UPDATES TO THE PEDIATRIC USE SECTION BASED ON THE PEDIATRIC STUDY REPORT ENTITLED, "A PHASE II PILOT TRIAL OF BORTEZOMIB IN COMBINATION WITH INTENSIVE RE-INDUCTION THERAPY IN CHILDREN WITH RELAPSED ACUTE LYMPHOBLASTIC LYMPHOMA (LL) "
- M-166 UPDATE TO LABELING WITH WEEK 48 RESULTS FROM VIKING-4 IN ANTIRETROVIRAL THERAPY (ART) - EXPERIENCED INTEGRASE STRAND TRANSFER INHIBITOR (INSTI) - RESISTANT SUBJECTS
- M-167 APPROVED FOR REVISIONS TO THE LABELING BASED ON THE CLINICAL STUDY ENTITLED "BRONCHOPULMONARY DYSPLASIA (BPD) IN PRETERM INFANTS REQUIRING MECHANICAL VENTILATION OR POSITIVE PRESSURE SUPPORT ON DAYS 5 TO 14 AFTER BIRTH".
- M-168 INFORMATION ADDED TO THE CLINICAL STUDIES SECTION OF THE LABELING REGARDING THE RE-NOVATE AND RE-NOVATE LL STUDIES (PROPHYLAXIS OF DEEP VEIN THROMBOSIS AND PULMONARY EMBOLISM FOLLOWING HIP REPLACEMENT SURGERY)
- M-169 UPDATES TO LABELING DESCRIBING RESPONSE TO A REPEAT COURSE OF PICATO GEL 0.015% ON THE FACE OR SCALP IF AN INCOMPLETE RESPONSE IS OBSERVED AT A FOLLOW-UP EXAMINATION.
- M-170 INFORMATION ADDED TO THE CLINICAL STUDIES SECTION REGARDING USE FOR THE TREATMENT OF PATIENTS WITH UNRESECTABLE OR METASTATIC MELANOMA WITH BRAF V600E OR V600K MUTATIONS AS DETECTED BY AN FDA-APPROVED TEST
- M-171 UPDATES TO LABELING WITH RESULTS TO THE TIGER CLINICAL TRIAL
- M-172 UPDATES TO THE CLINICAL TRIALS SECTION OF THE LABELING TO INCLUDE RESULTS OF STUDIES PERFORMED TO EVALUATE THE BENEFIT OF ADDING INCRUSE ELLIPTA TO PATIENTS WHO ARE ON BACKGROUND THERAPY WITH BREO ELLIPTA AND ADVAIR DISKUS
- M-173 INFORMATION ADDED TO THE CLINICAL STUDIES SECTION OF THE LABELING DESCRIBING THE EFFECTS OF STIOLTO RESPIMAT ON COPD PATIENTS
- M-174 INFORMATION ADDED TO CLINICAL STUDIES SECTION OF THE LABELING REGARDING INITIAL COMBINATION THERAPY OF EMPAGLIFLOZIN WITH METFORMIN
- M-175 INFORMATION ADDED TO THE LABELING DESCRIBING SAVOR, A PHASE IV TRIAL EVALUATING THE EFFECT OF SAXAGLIPTIN ON THE INCIDENCE OF CARDIOVASCULAR DEATH, MYOCARDIAL INFARCTION OR ISCHAEMIC STROKE IN PATIENTS WITH TYPE 2 DIABETES
- M-176 INFORMATION ADDED TO THE LABELING DESCRIBING TRIAL NN2211-3916, A TRIAL EVALUATING THE SAFETY AND EFFICACY OF LIRAGLUTIDE IN SUBJECTS WITH TYPE 2 DIABETES AND MODERATE RENAL IMPAIRMENT
- M-177 INFORMATION ADDED TO THE LABELING DESCRIBING EXAMINE, A TRIAL EVALUATING CARDIOVASCULAR ISCHEMIC RISKS ASSOCIATED WITH ALOGLIPTIN USE IN PATIENTS WITH TYPE 2 DIABETES AT HIGH RISK OF ISCHEMIC CARDIOVASCULAR DISEASE
- M-178 INFORMATION ADDED TO THE LABELING REGARDING MAINTENANCE OF REMISSION IN CROHN'S DISEASE IN PEDIATRIC PATIENTS
- M-179 UPDATES TO THE PRODUCT LABELING WITH STUDY REPORTS FROM THE OPTIMIST-1 AND OPTIMIST-2 CLINICAL TRIALS
- M-180 INFORMATION ADDED TO THE LABELING REGARDING THE ADDITION OF MAINTENANCE

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY MISCELLANEOUS**

TREATMENT IN PATIENTS WITH SCHIZOPHRENIA

- M-181 UPDATE TO THE DOSAGE AND ADMINISTRATION, PATIENT SELECTION (2.1), SECTION OF THE PACKAGE INSERT TO INCLUDE THE USE OF AN FDA-APPROVED PLASMA TEST FOR THE IDENTIFICATION OF EGFR EXON 19 DELETION OR EXON 21 (L858R) SUBSTITUTION MUTATIONS
- M-182 UPDATES TO THE PRODUCT LABELING BASED ON THE RESULTS OF STUDY H7T-MC-TADO TITLED, "A PHASE 3 DOUBLE-BLIND, RANDOMIZED, MULTICENTER, EFFICACY AND SAFETY STUDY OF PRASUGREL COMPARED TO PLACEBO IN PEDIATRIC PATIENTS WITH SICKLE CELL DISEASE"
- M-183 CHANGES TO THE DOSAGE AND ADMINISTRATION AND CLINICAL STUDIES SECTIONS OF THE LABELING TO SUPPORT THE REDUCE-TO-QUIT PARADIGM
- M-184 UPDATES MADE TO THE LABELING TO INCLUDE INFORMATION FROM STUDY MO25743 ON THE ANTI-TUMOR ACTIVITY OF VEMURAFENIB IN THE TREATMENT OF PATIENTS WITH BRAF V600E MUTATION-POSITIVE MELANOMA WITH BRAIN METASTASES
- M-185 UPDATES TO THE LABELING TO INCLUDE RESULTS OF A TRIAL TO EVALUATE THE SAFETY OF MOXIFLOXACIN IN PEDIATRIC PATIENTS WITH COMPLICATED INTRA-ABDOMINAL INFECTIONS
- M-186 UPDATES TO THE PRODUCT INFORMATION REGARDING MAINTENANCE TREATMENT OF SCHIZOPHRENIA IN ADULTS BASED UPON THE RESULTS FROM STUDY 331-10-232
- M-187 ADDITION OF CLINICAL INFORMATION OBTAINED FROM A PEDIATRIC TRIAL TO SECTION 8.4 OF THE LABELING
- M-188 PROVIDES FOR DATA SUPPORTING THE SAFETY AND EFFECTIVENESS FOR THE MAINTENANCE TREATMENT OF MODERATE TO SEVERE BINGE EATING DISORDER (BED)
- M-189 LABELING DESCRIBING THE EXPECTED REDUCTION OF ABUSE OF SINGLE-ENTITY MORPHINE BY THE INTRANASAL ROUTE OF ADMINISTRATION DUE TO PHYSICOCHEMICAL PROPERTIES
- M-190 INFORMATION ADDED TO THE CLINICAL STUDIES SECTION OF THE LABELING REGARDING THE LACK OF EFFICACY OF TARCEVA IN MAINTENANCE TREATMENT OF PATIENTS WITHOUT EGFR MUTATIONS
- M-191 ADDITION OF DATA BASED ON PEDIATRIC STUDIES TO FULFILL THE POSTMARKETING REQUIREMENT 1857-2
- M-192 PROVIDES FOR DATA EVALUATING THE NEUROPSYCHIATRIC SAFETY AND EFFICACY OF VARENICLINE FOR SMOKING CESSATION IN SUBJECTS WITH AND WITHOUT A HISTORY OF PSYCHIATRIC DISORDERS
- M-193 INFORMATION ADDED TO THE LABELING REGARDING A 15-WEEK, RANDOMIZED, DOUBLE-BLIND, PARALLEL-GROUP, PLACEBO-CONTROLLED FLEXIBLE-DOSE SAFETY AND EFFICACY STUDY OF PREGABALIN IN ADOLESCENTS (12 THROUGH 17 YEARS OLD) WITH FIBROMYALGIA
- M-194 INFORMATION ADDED TO THE LABELING REGARDING USE OF REGADENOSON ADMINISTRATION FOLLOWING AN INADEQUATE EXERCISE STRESS TEST AS COMPARED TO REGADENOSON ALONE
- M-195 REVISIONS TO THE PEDIATRIC USE SECTION OF THE LABELING REFLECTING LACK OF EFFICACY FOR IRRITABILITY ASSOCIATED WITH AUTISTIC DISORDER IN PEDIATRIC PATIENTS AGES 6-17
- M-196 REVISIONS TO THE PACKAGE INSERT BASED ON DATA FROM A RANDOMIZED, PLACEBO CONTROLLED, MULTICENTER STUDY OF INTRAVENOUS ACETAMINOPHEN FOR THE TREATMENT OF ACUTE PAIN IN PEDIATRIC PATIENTS TO FULFILL THE POST-MARKETING REQUIREMENT 1704-1
- M-197 NEW CLINICAL DATA ADDED TO THE PRESCRIBING INFORMATION REGARDING CANAGLIFLOZIN ADD-ON COMBINATION THERAPY WITH METFORMIN AND A DIPEPTIDYL-PEPTIDASE-4 INHIBITOR
- M-198 PACKAGE INSERT UPDATED WITH RESULTS FROM STUDY CV181168, A MULTICENTER, RANDOMIZED, DOUBLE-BLIND, PLACEBO-CONTROLLED, PARALLEL GROUP, PHASE 3 TRIAL TO EVALUATE THE SAFETY AND EFFICACY OF SAXAGLIPTIN ADDED TO DAPAGLIFLOZIN AND METFORMIN
- M-199 INFORMATION ADDED TO LABELING REGARDING THE TREATMENT OF PATIENTS WITH ALK-POSITIVE NON-SMALL CELL LUNG CANCER (NSCLC) WHO HAD NOT RECEIVED PRIOR SYSTEMIC THERAPY FOR METASTATIC DISEASE.
- M-200 CLINICAL INFORMATION ADDED TO THE USE IN SPECIFIC POPULATIONS SECTION OF THE LABELING.
- M-201 REVISIONS TO THE PACKAGE INSERT BASED ON DATA FROM AN OPEN LABEL, MULTI-CENTER STUDY OF CABAZITAXEL IN PEDIATRIC PATIENTS WITH REFRACTORY SOLID TUMORS INCLUDING TUMORS OF THE CENTRAL NERVOUS SYSTEM.
- M-202 INCLUSION OF DATA FROM THE SUMMIT STUDY FOR BREO ELLIPTA (FLUTICASONE FUROATE/VILANTEROL TRIFENATATE) INHALATION POWDER IN THE PACKAGE INSERT.

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY MISCELLANEOUS**

- M-203 PROVIDES FOR REVISIONS TO THE PACKAGE INSERT TO REFLECT RESULTS OF TWO POSTMARKETING REQUIREMENT STUDIES ROP111662 AND ROP111569
- M-204 CLINICAL INFORMATION ADDED TO THE PACKAGE INSERT REGARDING USE OF ATORVASTATIN IN CHILDREN AND ADOLESCENTS AGES 10-17 WITH HETEROZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA (HEFH)
- M-205 INFORMATION ADDED TO THE LABELING REGARDING RANDOMIZED, MULTICENTER, DOUBLE-BLIND, PLACEBO-CONTROLLED STUDIES ON PATIENTS WITH SEVERE RENAL IMPAIRMENT
- M-206 INFORMATION ADDED TO LABELING REGARDING 48 WEEK EFFICACY, RESISTANCE AND SAFETY DATA ON VIROLOGICALLY SUPPRESSED HIV-1 INFECTED ADULTS SWITCHING FROM COMPLERA TO ODEFSEY
- M-207 INFORMATION ADDED TO LABELING REGARDING 48 WEEK EFFICACY, RESISTANCE AND SAFETY DATA ON VIROLOGICALLY SUPPRESSED HIV-1 INFECTED ADULTS SWITCHING FROM ATRIPLA TO ODEFSEY
- M-208 INFORMATION ADDED TO THE LABELING TO INCLUDE RESULTS OF A RANDOMIZED, DOUBLE-BLIND, PLACEBO-CONTROLLED STUDY IN PATIENTS WITH SEVERE COPD ASSOCIATED WITH CHRONIC BRONCHITIS AND A HISTORY OF EXACERBATIONS
- M-209 INFORMATION ADDED TO THE LABELING REGARDING CABAZITAXEL AT 20 MG/M2 BASED ON THE RESULTS OF THE PROSELICA STUDY
- M-210 INFORMATION ADDED TO LABELING TO SUPPORT THE USE OF SYMBICORT TO REDUCE EXACERBATIONS IN PATIENTS WITH CHRONIC OBSTRUCTIVE PULMONARY DISEASE (COPD)
- M-211 PROVIDES FOR LABELING CHANGES REGARDING THE USE OF DAPTOMYCIN IN THE PEDIATRIC POPULATION FOR STAPHYLOCOCCUS AUREUS BACTEREMIA (SAB) BASED ON RESULTS OF A TRIAL IN PEDIATRIC PATIENTS 1 TO 17 YEARS OF AGE
- M-212 INFORMATION ADDED TO THE LABELING REGARDING THE SAFETY AND EFFICACY OF DAPAGLIFLOZIN IN PATIENTS WITH TYPE 2 DIABETES WHO HAVE INADEQUATE GLYCEMIC CONTROL ON A BACKGROUND COMBINATION OF METFORMIN AND EXENATIDE EXTENDED RELEASE
- M-213 INFORMATION ADDED TO THE LABELING TO INCLUDE THE EFFICACY AND SAFETY OF CARIPRAZINE RELATIVE TO PLACEBO IN THE PREVENTION OF RELAPSE OF SYMPTOMS IN PATIENTS WITH SCHIZOPHRENIA
- M-214 INFORMATION ADDED TO THE CLINICAL TRIALS SECTION OF THE LABELING REGARDING A POSTMARKETING SAFETY AND EFFICACY STUDY EVALUATING THE RISK OF SERIOUS ASTHMA-RELATED EVENTS
- M-215 INFORMATION ADDED TO THE LABELING REGARDING THE COMPARISON OF PALIPERIDONE PALMITATE COMPARED WITH ORAL ANTIPSYCHOTIC TREATMENT IN DELAYING TIME TO TREATMENT FAILURE IN ADULTS WITH SCHIZOPHRENIA WHO HAVE BEEN INCARCERATED
- M-216 UPDATE THE PRESCRIBING INFORMATION AND PATIENT LABELING WITH FINDINGS FROM STUDY RP103-08 CONDUCTED IN TREATMENT-NAIVE NEPHROPATHIC CYSTINOSIS PATIENTS TO EXPAND THE INDICATED POPULATION TO PATIENTS 1 YEAR AND OLDER
- M-217 INCORPORATION OF THE LABELING REVISIONS PROVIDED FOR IN NDA 022253/S-039 AND NDA 022255/S-022 INTO THE LACOSAMIDE INJECTION LABELING
- M-218 ADDITIONAL INFORMATION ADDED TO THE PEDIATRIC USE SECTION OF THE LABELING REGARDING A NEW CLINICAL TRIAL IN PATIENTS AGED 6 THROUGH 11 YEARS (TRIAL 4)
- M-219 INFORMATION ADDED TO THE PEDIATRIC USE SECTION OF THE LABELING REGARDING A NEW CLINICAL TRIAL IN PATIENTS 7 TO 14 YEARS OF AGE WITH DUCHENNE MUSCULAR DYSTROPHY
- M-220 ADDITIONAL INFORMATION ADDED TO THE LABELING FROM STUDY PC B308/13 REGARDING THE USE OF BLUE LIGHT CYSTOSCOPY WITH CYSVIEW AS AN ADJUNCT TO WHITE LIGHT CYSTOSCOPY
- M-221 DRUG FACTS LABELING CHANGES UNDER THE DIRECTIONS HEADING TO REVISE THE STATED PREPARATION TIME OF A DRY SITE FROM 120 SECONDS SCRUBBING AND 90 SECONDS DRYING TO 30 SECONDS SCRUBBING AND 30 SECONDS DRYING
- M-222 ADDITION OF DATA BASED ON THE ASSESSMENT OF SAFETY AND EFFICACY IN PEDIATRIC PATIENTS WITH MAJOR DEPRESSIVE DISORDER TO FULFILL POSTMARKETING STUDY REQUIREMENT 1229-1
- M-223 INFORMATION ADDED TO SECTION 8.1 OF THE LABELING REGARDING PREGNANT PATIENTS WHO ARE ALREADY ON A STABLE RILPIVIRINE REGIMEN PRIOR TO PREGNANCY AND WHO ARE VIROLOGICALLY SUPPRESSED (HIV-1 RNA LESS THAN 50 COPIES/ML)
- M-224 INFORMATION ADDED TO THE LABELING REGARDING THE SAFETY AND EFFICACY OF EXENATIDE EXTENDED RELEASE AS ADD-ON IN PATIENTS WITH TYPE 2 DIABETES WHO HAVE INADEQUATE GLYCEMIC CONTROL ON BASAL INSULIN GLARGINE WITH OR WITHOUT METFORMIN

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY MISCELLANEOUS**

- M-225 REVISIONS TO SECTION 8.4 OF THE PRESCRIBING INFORMATION TO INCLUDE A SAFETY AND EFFICACY STUDY IN PEDIATRIC PATIENTS AGES ≥ 6 YEARS TO < 18 YEARS WITH CHRONIC IDIOPATHIC CONSTIPATION
- M-226 CHANGES TO THE LABELING BASED ON RESULTS FROM A CONTROLLED CLINICAL TRIAL IN PATIENTS WITH LATER-ONSET SPINAL MUSCULAR ATROPHY
- M-227 ADDITION TO THE CLINICAL STUDIES SECTION OF THE LABELING WITH THE SUBSECTION ENTITLED DIGIT SYMBOL SUBSTITUTION TEST IN MAJOR DEPRESSIVE DISORDER
- M-228 INFORMATION ADDED TO THE PACKAGE INSERT REGARDING THE REVISION OF THE MONOTHERAPY INDICATION OF VENETOCLAX
- M-229 REVISED LABELING TO INCORPORATE THE PEDIATRIC USE OF LOTEHPREDNOL ETABONATE GEL IN PATIENTS FOR THE TREATMENT OF POSTOPERATIVE INFLAMMATION FOLLOWING OCULAR SURGERY
- M-230 REVISIONS TO THE GLECAPREVIR/PIBRENTASVIR COMBINATION PRODUCT PRESCRIBING INFORMATION TO INCLUDE SAFETY AND EFFICACY DATA FROM THE HCV/HIV-1 COINFECTION STUDY M14-730 AND FROM THE LIVER AND RENAL TRANSPLANT STUDY M13-596
- M-231 REVISIONS TO THE USE IN SPECIFIC POPULATIONS SECTION (SECTION 8.3) OF THE PACKAGE INSERT WITH THE RESULTS OF CLINICAL TRIAL WV25651, CONDUCTED TO EVALUATE THE EFFECT OF VALGANCICLOVIR ON SPERMATOGENESIS AND TO FULFILL PMR 1670-3
- M-232 INFORMATION ADDED TO SECTION 8.4 OF THE LABELING TO DESCRIBE THE RESULTS FROM PEDIATRIC STUDIES
- M-233 INFORMATION ADDED TO THE LABELING TO DESCRIBE FIXED-DOSE COMBINATION OF TIOTROPIUM BROMIDE AND OLODATEROL TO INCLUDE REDUCTION OF COPD EXACERBATIONS
- M-234 UPDATE TO THE PRESCRIBING INFORMATION FOR VORTIOXETINE ON TREATMENT-EMERGENT SEXUAL DYSFUNCTION COMPARING VORTIOXETINE AND SSRIS
- M-235 INFORMATION ADDED TO SECTION 14 OF THE LABELING TO DESCRIBE STUDY LAP016A2307 TO FULFILL POSTMARKETING STUDY REQUIREMENT 1586-1
- M-236 INFORMATION ADDED TO THE PRESCRIBING INFORMATION TO INCLUDE EFFICACY AND SAFETY DATA FROM A STUDY IN PATIENTS WITH TREATMENT NAIVE CLL/SLL TREATED WITH IBRUTINIB IN COMBINATION WITH OBINUTUZUMAB OR CHLORAMBUCIL IN COMBINATION WITH OBINUTUZUMAB
- M-237 INFORMATION ADDED TO LABELING TO DESCRIBE A STUDY TO EVALUATE THE SAFETY AND EFFICACY OF VARENICLINE FOR SMOKING CESSATION IN ADOLESCENT SMOKERS
- M-238 INFORMATION ADDED TO THE PRESCRIBING INFORMATION TO REFLECT THAT NO DOSE ADJUSTMENT IS NEEDED FOR PATIENTS WITH AN ESTIMATED GLOMERULAR FILTRATION RATE (EGFR) OF 45 ML/MIN/1.73 M² OR GREATER AS SUPPORTED BY CLINICAL STUDY REPORT
- M-239 INFORMATION ADDED TO THE PEDIATRIC USE SECTION OF THE LABELING REGARDING A TRIAL CONDUCTED IN TREATMENT NAIVE PEDIATRIC PATIENTS, AGES 2 YEARS TO < 18 YEARS WITH TRANSFUSIONAL IRON OVERLOAD
- M-240 INFORMATION ADDED TO LABELING REGARDING A RANDOMIZED, PLACEBO-CONTROLLED CLINICAL TRIAL TO EVALUATE CARDIOVASCULAR OUTCOMES AFTER TREATMENT WITH EXENATIDE ONCE WEEKLY IN PATIENTS WITH TYPE 2 DIABETES MELLITES
- M-241 INFORMATION ADDED TO THE LABELING FOR SAFETY & EFFICACY STUDY ENTITLED, A MULTICENTER, RANDOMIZED, DOUBLE-BLIND, PLACEBO CONTROLLED CLINICAL TRIAL OF DEFERASIROX IN PATIENTS WITH MYELODYSPLASTIC SYNDROMES (LOW/INT-1 RISK) & TRANSFUSIONAL IRON OVERLOAD
- M-242 INFORMATION ADDED TO THE LABELING REGARDING THE EFFICACY AND SAFETY OF INSULIN DEGLUDEC/LIRAGLUTIDE VS INSULIN GLARGINE IN PTS W/ TYPE 2 DIABETES INADEQUATELY CONTROLLED ON SGLT2I WITH OR WITHOUT ORAL ANTIDIABETIC THERAPIES
- M-243 INFORMATION ADDED TO LABELING FROM PROSPECTIVE, RANDOMIZED, OPEN-LABEL, BLIND EVALUATOR (PROBE) STUDY EVALUATING THE EFFICACY AND SAFETY OF LOW MOLECULAR WEIGHT HEPARIN/EDOXABAN VERSUS DALTEPARIN IN VENOUS THROMBOEMBOLISM ASSOCIATED WITH CANCER
- M-244 INFORMATION ADDED TO THE LABELING REGARDING EFFICACY AND SAFETY OF THE CONTINUATION OF SITAGLIPTIN COMPARED WITH THE WITHDRAWAL OF SITAGLIPTIN DURING INITIATION AND TITRATION OF INSULIN GLARGINE IN SUBJECTS WITH TYPE 2 DIABETES MELLITUS
- M-245 ADDITIONAL INFORMATION ADDED TO THE LABELING BASED ON SAFETY AND EFFICACY DATA FROM THE IMPACT TRIAL
- M-246 ADDITION OF STUDY BR117277, A NON-RANDOMIZED, OPEN-LABEL, MULTI-CENTER, MULTI-COHORT TRIAL OF DABRAFENIB PLUS TRAMETINIB IN SUBJECTS WITH BRAF MUTATION-

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY MISCELLANEOUS**

POSITIVE MELANOMA THAT HAS METASTASIZED TO THE BRAIN

M-247 REVISIONS TO THE LABELING REGARDING CONTINUOUS SUBCUTANEOUS INSULIN INFUSION AS A CONDITION OF USE FOR INSULIN ASPART

M-248 INFORMATION ADDED TO THE LABELING TO DESCRIBE A TRIAL EVALUATING A LOWER DOSE THAN THOSE APPROVED FOR PEDIATRIC PATIENTS 13 TO 17 YEARS OF AGE

M-249 INFORMATION ADDED TO THE LABELING TO DESCRIBE STUDY LVM-MD-15 TO FULFILL POSTMARKETING COMMITMENT 1943-4

M-250 REVISIONS TO THE PEDIATRIC USE SECTION TO INCLUDE AN OPEN-LABEL CLINICAL TRIAL TO FULFILL PMR 1655-1

M-251 INFORMATION ADDED TO THE CLINICAL PHARMACOLOGY SECTION REGARDING INFLUENZA VIRUS RESISTANCE TO OSELTAMIVIR IN IMMUNOCOMPROMISED PATIENTS

M-252 ADDITION OF INFORMATION TO CLINICAL STUDIES SECTION REGARDING CARDIOVASCULAR OUTCOME

M-253 INFORMATION ADDED TO THE LABELING TO DESCRIBE STUDY P061, A RANDOMIZED, PLACEBO-CONTROLLED, PARALLEL GROUP, MULTI-SITE, DOUBLE-BLIND STUDY TO EVALUATE SAFETY AND EFFICACY OF SUVOREXANT FOR THE TREATMENT OF INSOMNIA IN SUBJECTS WITH ALZHEIMERS DISEASE

M-254 INFORMATION ADDED TO THE LABELING REGARDING PEDIATRIC PATIENTS AGES 7 TO 17 YEARS OF AGE WITH MAJOR DEPRESSIVE DISORDER

M-255 INFORMATION ADDED TO THE LABELING TO DESCRIBE STUDY GS-US-320-4018 IN VIROLOGICALLY SUPPRESSED ADULTS W/ CHRONIC HEP B INFECTION WHO SWITCHED FROM TENOFOVIR DISOPROXIL FUMARATE TO TENOFOVIR ALAFENAMIDE

M-256 INFORMATION ADDED TO THE CLINICAL STUDIES SECTION TO FULFILL A POST-MARKETING REQUIREMENT

M-257 INFORMATION ADDED TO THE CLINICAL STUDIES SECTION OF THE LABELING REGARDING THE USE OF PLAQUE PSORIASIS OF THE SCALP

M-258 INFORMATION ADDED TO THE LABELING TO DESCRIBE CARMELINA TRIAL TO FULFILL POSTMARKETING COMMITMENT 1766-4

M-259 INFORMATION ADDED TO THE LABELING REGARDING SAFETY AND EFFICACY IN SUBJECTS WITH HCV SUBTYPE 3B INFECTION

M-260 INFORMATION ADDED TO THE LABELING DESCRIBING A RANDOMIZED, OPEN-LABEL STUDY THAT EXAMINED THE CONCOMITANT USE OF DIMETHYL FUMARATE AND SEVERAL NON-LIVE VACCINES IN ADULTS 27-55 YEARS OF AGE WITH RELAPSING FORMS OF MULTIPLE SCLEROSIS

M-261 ADDITIONAL INFORMATION ADDED TO THE LABELING REGARDING THE USE IN PATIENTS ON CHRONIC HEMODIALYSIS

M-262 REVISIONS TO THE USE IN SPECIFIC POPULATIONS SECTION OF THE PACKAGE INSERT TO INCLUDE THE RESULT OF STUDY P146 TO FULFILL THE REQUIREMENTS OF PMR 3003-4

M-263 REVISIONS TO THE LABELING TO INCLUDE RESULTS FROM CLINICAL STUDY ICL670E2419 (THETIS TRIAL) TO SUPPORT PMR 3342-2 AND 3342-3

M-264 INFORMATION ADDED TO THE LABELING DESCRIBING A PHASE 2, MULTICENTER, OPEN-LABEL STUDY TO EVALUATE THE SAFETY/EFFICACY OF SOFOSBUVIR/VELPATASVIR IN SUBJECTS WITH CHRONIC HCV INFECTION WHO HAVE RECEIVED A LIVER TRANSPLANT

M-265 REVISIONS TO THE LABELING TO INCLUDE RESULTS FROM CLINICAL STUDY M15-656 (VIALE-A) AND M16-043 (VIALE-C) TO SUPPORT PMR 3545-1 AND PMR 3545-2

M-266 INFORMATION ADDED TO THE LABELING TO DESCRIBE STUDY GS-US-320-4035 IN VIROLOGICALLY SUPPRESSED ADULTS W/ CHRONIC HEP B INFECTION WHO SWITCHED FROM TENOFOVIR DISOPROXIL FUMARATE TO TENOFOVIR ALFAENAMIDE

M-267 INFORMATION ADDED TO THE LABELING REGARDING THE RESULT OF STUDY LUAA21004-402

M-268 ADDITION OF INFORMATION TO THE LABEL REGARDING A CLEAR PRODUCT PRESENTATION AND 26 ML VOLUME PRODUCTS

M-269 REVISIONS TO THE LABELING TO INCLUDE RESULTS FROM CLINICAL STUDY 309/KEYNOTE-775 TO SUPPORT PMR 3696-1 AND 3700-1

M-270 INFORMATION ADDED TO CLINICAL PHARMACOLOGY SECTION

M-271 REVISIONS TO THE LABELING TO INCLUDE RESULTS FROM CLINICAL STUDY HRA2914-5016

M-272 REVISIONS TO THE LABELING TO INCLUDE RESULTS FROM CLINICAL STUDY E7080-G000-211 TO SUPPORT PMR 2865-1

M-273 REVISION TO THE LABELING TO INCLUDE RESULTS FROM CLINICAL STUDY 207966 ATLAS-2M

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY MISCELLANEOUS**

M-274 REVISIONS TO THE LABELING TO INCLUDE RESULTS FROM CLINICAL STUDY CBAF312A2130

M-275 REVISION TO THE LABELING TO INCLUDE RESULTS FROM CLINICAL STUDY, MK-8835-004/B1521021, VERTIS CV

M-276 REVISIONS TO THE PEDIATRIC USE SUBSECTION OF LABELING TO INCLUDE THE RESULTS FROM CLINICAL STUDY 1200.120, CONDUCTED TO FULFILL A PEDIATRIC WRITTEN REQUEST

M-277 UPDATES THE US PRESCRIBING INFORMATION WITH CLINICAL DATA REGARDING THE USE OF SOFOSBUVIR AND VELPATASVIR FOR THE TREATMENT OF CHRONIC HCV GENOTYPE 1, 2, 3, 4, 5, OR 6 INFECTION IN PEOPLE WHO INJECT DRUGS (PWID), INCLUDING THOSE ON MEDICATION-ASSISTED TREATMENT (MAT) FOR OPIOID USE DISORDER

M-278 INFORMATION ADDED TO SECTION 8.4 OF THE LABELING TO INCLUDE THE RESULT OF STUDY GO29665

M-279 INFORMATION ADDED TO THE LABELING TO DESCRIBE THE RESULTS OF FIGARO-DKD STUDY

M-280 REVISIONS TO THE PEDIATRIC USE SUBSECTION OF LABELING TO INCLUDE THE RESULTS FROM CLINICAL STUDIES E7389-G000-223 AND E7389-G000-213, CONDUCTED TO FULFILL A PEDIATRIC WRITTEN REQUEST

M-281 REVISIONS TO THE LABELING TO PROVIDE FOR THE EXPANSION OF THE USE OF STIRIPENTOL FOR THE TREATMENT OF SEIZURES ASSOCIATED WITH DRAVET SYNDROME IN PATIENTS TAKING CLOBAZAM TO INCLUDE PEDIATRIC PATIENTS WHO ARE 6 MONTHS TO LESS THAN 2 YEARS OF AGE AND WEIGHING 7 KG OR MORE

M-282 REVISIONS TO THE LABELING TO ADD THE RESULTS OF A CLINICAL STUDY (TA-303) IN PATIENTS WITH ED FOLLOWING BILATERAL NERVE-SPARING RADICAL PROSTATECTOMY

M-283 INFORMATION ADDED TO SECTION 8.4 OF THE LABELING TO INCLUDE THE RESULT OF STUDY HESTIA3

M-284 REVISIONS TO THE LABELING TO INCLUDE RESULTS FROM GALILEO TRIAL

M-285 REVISIONS TO SECTION 8.4 OF THE LABELING TO INCLUDE THE RESULTS OF STUDY INCB 18424-269

M-286 INFORMATION ADDED TO CLINICAL PHARMACOLOGY SECTION TO INCLUDE RESULTS FROM STUDY ORGN001-102

M-287 LABELING REGARDING NEW DOSING REGIMEN IN ADULTS WITH PULMONARY ARTERIAL HYPERTENSION (PAH) AND TREATMENT OF PAH IN PEDIATRIC PATIENTS (AGES 1-17)

M-288 INFORMATION ADDED TO THE LABELING TO DESCRIBE THE RESULTS OF STUDY APL2-308

M-289 INFORMATION ADDED TO THE LABELING TO DESCRIBE THE RESULTS OF MVT-601-035

M-290 INFORMATION ADDED TO SECTION 8.4 OF THE LABELING TO INCLUDE THE RESULT OF STUDY HZA114971

M-291 REVISIONS TO THE USE IN SPECIFIC POPULATIONS SECTION OF THE PACKAGE INSERT TO INCLUDE THE RESULT OF STUDY P145

M-292 REVISIONS TO THE LABELING TO DESCRIBE MODIFIED FORMULATION BASED ON RESULTS OF STUDIES EM-05-014624 AND EM-05-014815

M-293 REVISIONS TO THE LABELING TO INCLUDE RESULTS FROM CLINICAL STUDY E2006-A001-113

M-294 INFORMATION ADDED TO THE LABELING REGARDING THE RESULT OF STUDY 205860

M-295 REVISIONS TO THE PEDIATRIC USE SUBSECTION OF LABELING TO INCLUDE THE RESULTS FROM CLINICAL STUDY 1218-0091, CONDUCTED TO FULFILL A PEDIATRIC WRITTEN REQUEST

M-296 REVISIONS TO THE LABELING TO INCLUDE RESULTS FROM PROTOCOL 1218- 0091

M-297 LABELING CHANGES BASED ON RESULTS FROM CLINICAL STUDY MYK-461-017

M-298 LABELING REVISIONS RELATED TO STUDY D1699CC00001

M-299 CLINICAL STUDY INFORMATION ADDED TO THE LABEL ABOUT THE TREATMENT OF MODERATE TO SEVERE GENITAL PSORIASIS

M-300 REVISIONS TO THE PEDIATRIC USE SUBSECTION OF LABELING TO INCLUDE THE RESULTS FROM CLINICAL STUDY SYR-322-309, CONDUCTED IN RESPONSE TO A PEDIATRIC WRITTEN REQUEST

M-301 CLINICAL STUDY INFORMATION ADDED TO LABEL ABOUT THE TREATMENT OF CORONAVIRUS DISEASE 2019 (COVID-19) IN PATIENTS WITH SEVERELY REDUCED RENAL FUNCTION (ESTIMATED GLOMERULAR FILTRATION RATE, EGFR < 30 ML/MIN)

M-302 INFORMATION ADDED TO LABELING REGARDING OSTEOSARCOMA

M-303 INFORMATION ADDED TO THE CLINICAL STUDIES SECTION TO FULFILL A POST-MARKETING REQUIREMENT

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY MISCELLANEOUS**

M-304 INFORMATION ADDED TO SECTION 8.4 OF THE LABELING TO DESCRIBE THE RESULTS FROM STUDY LVM-MD-11 AND LVM-MD-14

M-305 REVISIONS TO THE LABELING TO INCLUDE INFORMATION FOR PREGNANT INDIVIDUALS

M-306 REVISIONS TO THE LABELING TO REFLECT THE RESULTS OF A CLINICAL STUDY TO FULFILL POST MARKETING REQUIREMENT 1988-001

M-307 INFORMATION ADDED TO THE CLINICAL PHARMACOLOGY SECTION TO INCLUDE RESULTS FROM A DRUG INTERACTION STUDY WITH GASTRIC ACID REDUCING AGENTS

M-308 INFORMATION ADDED TO THE LABELING TO DESCRIBE A CLINICAL STUDY

M-309 INFORMATION ADDED TO THE LABELING TO INCLUDE RESULTS FROM A STUDY TO FULFILL POSTMARKETING REQUIREMENT 3809-5

M-310 REVISIONS TO THE CLINICAL STUDIES SECTION TO REFLECT THE RESULTS OF A CLINICAL STUDY TO FULFILL POST-MARKETING REQUIREMENT 4150-1

M-311 REVISIONS TO THE LABELING TO REFLECT THE RESULTS OF STUDY LIBRETTO-431

M-312 REVISIONS TO THE LABELING TO REFLECT THE RESULTS OF STUDY LIBRETTO-531

ORPHAN DRUG EXCLUSIVITY

ODE-1 TO REDUCE CHRONIC DROOLING IN PATIENTS AGED 3 - 16 WITH NEUROLOGIC CONDITIONS ASSOCIATED WITH PROBLEM DROOLING (E.G. CEREBRAL PALSY)

ODE-2 FOR TREATMENT OF NON-INFECTIOUS UVEITIS AFFECTING THE POSTERIOR SEGMENT OF THE EYE

ODE-3 TO TREAT INFANTILE SPASMS

ODE-4 TREATMENT OF PATIENTS WITH SUBEPENDYMAL GIANT CELL ASTROCYTOMA (SEGA) ASSOCIATED WITH TUBEROUS SCLEROSIS WHO REQUIRE THERAPEUTIC INTERVENTION BUT ARE NOT CANDIDATES FOR CURATIVE SURGICAL RESECTION

ODE-5 FOR SEQUENTIAL USE FOR THE TREATMENT OF CYANIDE POISONING THAT IS JUDGED TO BE LIFE-THREATENING

ODE-6 FOR THE MANAGEMENT OF POSTHERPETIC NEURALGIA

ODE-7 TO REDUCE THE RISK OF PRETERM BIRTH IN WOMEN WITH SINGLETON PREGNANCY WHO HAVE A HISTORY OF SINGLETON SPONTANEOUS PRETERM BIRTH

ODE-8 TREATMENT OF SEVERE HYPERCALCEMIA IN PATIENTS WITH PRIMARY HYPERPARATHYROIDISM WHO ARE UNABLE TO UNDERGO PARATHYROIDECTOMY

ODE-9 TREATMENT OF ASYMPTOMATIC OR PROGRESSIVE MEDULLARY THYROID CANCER IN PATIENTS WITH UNRESECTABLE LOCALLY ADVANCED OR METASTATIC DISEASE

ODE-10 FOR USE IN COMBINATION CHEMOTHERAPY WITH 5-FLUOROURACIL IN THE PALLIATIVE TREATMENT OF PATIENTS WITH ADVANCED METASTATIC COLORECTAL CANCER

ODE-11 TREATMENT OF PROGRESSIVE NEUROENDOCRINE TUMORS OF PANCREATIC ORIGIN (PNET) IN PATIENTS WITH UNRESECTABLE, LOCALLY ADVANCED OR METASTATIC DISEASE

ODE-12 TREATMENT OF PERIPHERAL T-CELL LYMPHOMA (PTCL) IN PATIENTS WHO HAVE RECEIVED AT LEAST ONE PRIOR THERAPY

ODE-13 TREATMENT OF UNRESECTABLE OR METASTATIC MELANOMA WITH THE BRAF V600E MUTATION AS DETECTED BY AN FDA-APPROVED TEST

ODE-14 TREATMENT OF ACUTE ATTACKS OF HEREDITARY ANGIOEDEMA IN ADULTS 18 YEARS OF AGE AND OLDER

ODE-15 TREATMENT OF PATIENTS WITH LOCALLY ADVANCED OR METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) THAT IS ANAPLASTIC LYMPHOMA KINASE (ALK)-POSITIVE AS DETECTED BY AN FDA APPROVED TEST

ODE-16 TREATMENT OF PATIENTS WITH TRANSFUSIONAL IRON OVERLOAD DUE TO THALASSEMIA SYNDROMES WHEN CURRENT CHELATION THERAPY IS INADEQUATE

ODE-17 ADJUNCTIVE TREATMENT OF SEIZURES ASSOCIATED WITH LENNOX-GASTAUT SYNDROME IN PATIENTS 2 YEARS OF AGE OR OLDER

ODE-19 TREATMENT OF PATIENTS WITH INTERMEDIATE OR HIGH-RISK MYELOFIBROSIS, INCLUDING PRIMARY MYELOFIBROSIS, POST-POLYCYTHEMIA VERA MYELOFIBROSIS AND POST-ESSENTIAL THROMBOCYTHEMIA MYELOFIBROSIS

ODE-20 TREATMENT OF CYSTIC FIBROSIS (CF) IN PATIENTS AGE 6 YEARS AND OLDER WHO HAVE A

PATENT AND EXCLUSIVITY TERMS**ORPHAN DRUG EXCLUSIVITY**

- G551D MUTATION IN THE CFTR GENE.
- ODE-21 AN ADJUNCT TO AB EXTERNO GLAUCOMA SURGERY
- ODE-22 FOR THE CONTROL OF HYPERGLYCEMIA SECONDARY TO HYPERCORTISOLISM IN ADULT PATIENTS WITH ENDOGENOUS CUSHING'S SYNDROME WHO HAVE TYPE 2 DIABETES MELLITUS OR GLUCOSE INTOLERANCE AND HAVE FAILED SURGERY OR ARE NOT CANDIDATES FOR SURGERY
- ODE-23 ADVANCED SOFT TISSUE SARCOMA (STS) WHO HAVE RECEIVED PRIOR CHEMOTHERAPY
- ODE-24 TREATMENT OF ADULTS WITH RENAL ANGIOMYOLIPOMA AND TUBEROUS SCLEROSIS COMPLEX (TSC) NOT REQUIRING IMMEDIATE SURGERY
- ODE-25 MANAGEMENT OF POSTHERPETIC NEURALGIA IN ADULTS.
- ODE-26 TREATMENT OF ENDOGENOUS ANTERIOR UVEITIS
- ODE-27 TREATMENT OF PATIENTS WITH MULTIPLE MYELOMA WHO HAVE RECEIVED AT LEAST TWO PRIOR THERAPIES INCLUDING BORTEZOMIB AND AN IMMUNOMODULATORY AGENT AND HAVE DEMONSTRATED DISEASE PROGRESSION ON OR WITHIN 60 DAYS OF COMPLETION OF THE LAST THERAPY
- ODE-28 TREATMENT OF ADULT PATIENTS WITH PHILADELPHIA CHROMOSOME-NEGATIVE (PH-) ACUTE LYMPHOBLASTIC LEUKEMIA (ALL) IN SECOND OR GREATER RELAPSE OR WHOSE DISEASE HAS PROGRESSED FOLLOWING TWO OR MORE ANTI-LEUKEMIA THERAPIES
- ODE-29 LOCALIZATION OF LYMPH NODES DRAINING A PRIMARY TUMOR IN PATIENTS WITH MELANOMA WHEN USED WITH A HAND-HELD GAMMA COUNTER
- ODE-30 TREATMENT OF ADULT PATIENTS WITH CHRONIC, ACCELERATED OR BLAST PHASE PHILADELPHIA CHROMOSOME-POSITIVE (PH+) CHRONIC MYELOGENOUS LEUKEMIA (CML) WITH RESISTANCE, OR INTOLERANCE TO PRIOR THERAPY
- ODE-31 TREATMENT OF CORNEAL CYSTINE CRYSTAL ACCUMULATION IN PATIENTS WITH CYSTINOSIS
- ODE-32 TREATMENT OF ADULT PATIENTS WITH CHRONIC OR ACCELERATED PHASE CHRONIC MYELOID LEUKEMIA (CML) WITH RESISTANCE AND/OR INTOLERANCE TO TWO OR MORE TYROSINE KINASE INHIBITORS (TKI)
- ODE-33 TREATMENT OF PROGRESSIVE, METASTATIC MEDULLARY THYROID CANCER (MTC)
- ODE-34 TREATMENT OF ADULT PATIENTS WITH CUSHING'S DISEASE FOR WHOM PITUITARY SURGERY IS NOT AN OPTION OR HAS NOT BEEN CURATIVE
- ODE-35 TREATMENT OF ADULT PATIENTS WITH PHILADELPHIA CHROMOSOME POSITIVE ACUTE LYMPHOBLASTIC LEUKEMIA (PH+ALL) THAT IS RESISTANT OR INTOLERANT TO PRIOR TYROSINE KINASE INHIBITOR THERAPY.
- ODE-36 ADJUNCT TO A LOW-FAT DIET AND OTHER LIPID-LOWERING TREATMENTS, INCLUDING LDL APHERESIS WHERE AVAILABLE, TO REDUCE LDL-C, TC, APOLIPOPROTEIN B, & NON-HDL-C IN PATIENTS WITH HOMOZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA
- ODE-37 TREATMENT OF ADULT PATIENTS WITH SHORT BOWEL SYNDROME (SBS) WHO ARE DEPENDENT ON PARENTERAL SUPPORT
- ODE-38 PART OF COMBINATION THERAPY IN ADULTS (GREATER THAN OR EQUAL TO 18 YEARS) WITH PULMONARY MULTI-DRUG RESISTANT TUBERCULOSIS (MDR-TB)
- ODE-39 TREATMENT OF CHRONIC IRON OVERLOAD IN PATIENTS 10 YRS. & OLDER WITH NON-TRANSFUSION DEPENDENT THALASSEMIA (NTDT) SYNDROMES AND WITH A LIVER IRON CONCENTRATION OF AT LEAST 5 MG OF IRON PER GRAM OF LIVER DRY WEIGHT & SERUM FERRITIN GREATER THAN 300 MCG/L.
- ODE-40 TREATMENT OF PEDIATRIC PATIENTS WITH NEWLY DIAGNOSED PHILADELPHIA CHROMOSOME-POSITIVE ACUTE LYMPHOBLASTIC LEUKEMIA (PH+ ALL) IN COMBINATION WITH CHEMOTHERAPY, APPROVED UNDER NDA #21588/S-037
- ODE-41 ADJUNCT TO LIPID-LOWERING MEDICATIONS AND DIET TO REDUCE LDL-C, APOLIPOPROTEIN B (APO B), TOTAL CHOLESTEROL (TC), AND NON-HIGH DENSITY LIPOPROTEIN-CHOLESTEROL (NON-HDL-C) IN PATIENTS WITH HOMOZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA (HOFH)
- ODE-42 USE AS A NITROGEN-BINDING ADJUNCTIVE THERAPY FOR CHRONIC MGMT OF ADULT AND PEDIATRIC PATIENTS AT LEAST 2 YRS WITH UREA CYCLE DISORDERS THAT CANNOT BE MANAGED BY DIETARY PROTEIN RESTRICTION AND/OR AMINO ACID SUPPLEMENTATION ALONE
- ODE-43 TREATMENT OF PATIENTS WITH MULTIPLE MYELOMA WHO HAVE RECEIVED AT LEAST TWO PRIOR THERAPIES INCLUDING LENALIDOMIDE AND BORTEZOMIB AND HAVE DEMONSTRATED DISEASE PROGRESSION ON OR WITHIN 60 DAYS OF COMPLETION OF THE LAST THERAPY.
- ODE-44 TREATMENT OF PATIENTS WITH LOCALLY ADVANCED, UNRESECTABLE OR METASTATIC GASTROINTESTINAL STROMAL TUMOR (GIST) WHO HAVE BEEN PREVIOUSLY TREATED WITH IMATINIB MESYLATE AND SUNITINIB MALATE.

PATENT AND EXCLUSIVITY TERMS**ORPHAN DRUG EXCLUSIVITY**

- ODE-45 MANAGEMENT OF NEPHROPATHIC CYSTINOSIS IN ADULTS AND CHILDREN AGES 6 YEARS AND OLDER.
- ODE-46 IMPROVEMENT OF NEUROLOGICAL OUTCOME BY REDUCING THE INCIDENCE AND SEVERITY OF ISCHEMIC DEFICITS IN ADULT PATIENTS WITH SUBARACHNOID HEMORRHAGE FROM RUPTURED INTRACRANIAL BERRY ANEURYSMS REGARDLESS OF THEIR POST-ICTUS NEUROLOGICAL CONDITION
- ODE-47 TREATMENT OF PATIENTS WITH UNRESECTABLE OR METASTATIC MELANOMA WITH BRAF V600E MUTATION AS DETECTED BY AN FDA APPROVED TEST.
- ODE-48 TREATMENT OF PATIENTS WITH UNRESECTABLE OR METASTATIC MELANOMA WITH BRAF V600E OR V600K MUTATIONS AS DETECTED BY AN FDA APPROVED TEST
- ODE-49 TREATMENT OF MANTLE CELL LYMPHOMA WHOSE DISEASE HAS RELAPSED OR PROGRESSED AFTER TWO PRIOR THERAPIES, ONE OF WHICH INCLUDED BORTEZOMIB
- ODE-50 FIRST-LINE TREATMENT OF PATIENTS WITH METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) WHOSE TUMORS HAVE EPIDERMAL GROWTH FACTOR RECEPTOR (EGFR) EXON 19 DELETIONS OR EXON 21 (L858R) SUBSTITUTION MUTATIONS AS DETECTED BY AN FDA-APPROVED TEST.
- ODE-51 TOPICAL TREATMENT OF STAGE 1A AND 1B MYCOSIS FUNGOIDES-TYPE CUTANEOUS T-CELL LYMPHOMA IN PATIENTS WHO HAVE RECEIVED PRIOR SKIN-DIRECTED THERAPY
- ODE-52 TREATMENT OF METASTATIC ADENOCARCINOMA OF THE PANCREAS AS FIRST-LINE TREATMENT, IN COMBINATION WITH GEMCITABINE.
- ODE-53 TREATMENT OF ADULTS WITH PULMONARY ARTERIAL HYPERTENSION (PAH) WHO GROUP 1, TO IMPROVE EXERCISE CAPACITY, WHO FUNCTIONAL CLASS AND TO DELAY CLINICAL WORSENING.
- ODE-54 TX OF PAH TO DELAY DISEASE PROGRESSION. DISEASE PROGRESSION INCLUDED: DEATH, INITIATION OF IV OR SC PROSTANOIDS, OR CLINICAL WORSENING OF PAH (DECREASED 6-MINUTE WALK DISTANCE, WORSENERD PAH SYMPTOMS AND NEED FOR ADDITIONAL PAH TREATMENT).
- ODE-55 TREATMENT OF PATIENTS WITH MANTLE CELL LYMPHOMA (MCL) WHO HAVE RECEIVED AT LEAST ONE PRIOR THERAPY
- ODE-56 TREATMENT OF PATIENTS WITH LOCALLY RECURRENT OR METASTATIC, PROGRESSIVE, DIFFERENTIATED THYROID CARCINOMA (DCT) THAT IS REFRACTORY TO RADIOACTIVE IODINE TREATMENT.
- ODE-57 TRAMETINIB IN COMBO WITH DABRAFENIB FOR TX. OF PTS WITH UNRESECTABLE OR METASTATIC MELANOMA WITH BRAF V600E OR V600K MUTATIONS AS DETECTED BY AN FDA-APPROVED TEST. THIS INDICATION IS BASED ON THE DEMONSTRATION OF DURABLE RESPONSE RATE
- ODE-58 DABRAFENIB IN COMBO WITH TRAMETINIB FOR TX. OF PTS WITH UNRESECTABLE OR METASTATIC MELANOMA WITH BRAF V600E OR V600K MUTATIONS AS DETECTED BY AN FDA-APPROVED TEST. THIS INDICATION IS BASED ON THE DEMONSTRATION OF DURABLE RESPONSE RATE
- ODE-59 TREATMENT OF NON-24-HOUR SLEEP-WAKE DISORDER
- ODE-60 TREATMENT OF PATIENTS WITH CHRONIC LYMPHOCYTIC LEUKEMIA (CLL) WHO HAVE RECEIVED AT LEAST ONE PRIOR THERAPY
- ODE-61 TREATMENT OF NEUROGENIC SYMPTOMATIC ORTHOSTATIC HYPOTENSION IN PATIENTS WITH PRIMARY AUTONOMIC FAILURE, DOPAMINE-BETA-HYDROXYLASE DEFICIENCY, AND NONDIABETIC AUTONOMIC NEUROPATHY
- ODE-62 TREATMENT OF PROLIFERATING INFANTILE HEMANGIOMA REQUIRING SYSTEMIC THERAPY.
- ODE-63 TREATMENT OF VISCERAL LEISHMANIASIS DUE TO LEISHMANIA DONOVANI; CUTANEOUS LEISHMANIASIS DUE TO LEISHMANIA BRAZILIENSIS, LEISHMANIA GUYANENSIS, AND LEISHMANIA PANAMENSIS; AND MUCOSAL LEISHMANIASIS DUE TO LEISHMANIA BRAZILIENSIS.
- ODE-64 SELECTIVE HEPATIC INTRA-ARTERIAL USE FOR IMAGING TUMORS IN ADULTS WITH KNOWN HEPATOCELLULAR CARCINOMA (HCC)
- ODE-65 TREATMENT OF PATIENTS WITH ACUTE LYMPHOBLASTIC LEUKEMIA AS PART OF A COMBINATION REGIMEN.
- ODE-66 TREATMENT OF PATIENTS WITH ANAPLASTIC LYMPHOMA KINASE (ALK)-POSITIVE METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) WHO HAVE PROGRESSED ON OR ARE INTOLERANT TO CRIZOTINIB.
- ODE-67 GUIDING SENTINEL LYMPH NODE BIOPSY, USING A HAND-HELD GAMMA COUNTER IN PATIENTS WITH CLINICALLY NODE NEGATIVE SQUAMOUS CELL CARCINOMA OF THE ORAL CAVITY
- ODE-68 TREATMENT OF PATIENTS WITH RELAPSED OR REFRACTORY PERIPHERAL T-CELL LYMPHOMA

PATENT AND EXCLUSIVITY TERMS**ORPHAN DRUG EXCLUSIVITY**

- ODE-69 TREATMENT OF MALIGNANT HYPERTHERMIA IN CONJUNCTION WITH APPROPRIATE SUPPORTIVE MEASURES AND FOR THE PREVENTION OF MALIGNANT HYPERTHERMIA IN PATIENTS AT HIGH RISK
- ODE-70 RELAPSED CLL, IN COMBO. WITH RITUXIMAB, IN PATIENTS FOR WHOM RITUXIMAB ALONE WOULD BE CONSIDERED APPROPRIATE THERAPY DUE TO OTHER CO-MORBIDITIES; AND RELAPSED SLL IN PATIENTS WHO HAVE RECEIVED AT LEAST 2 PRIOR SYSTEMIC THERAPIES
- ODE-71 RELAPSED FOLLICULAR B-CELL NON-HODGKIN LYMPHOMA (FL) IN PATIENTS WHO HAVE RECEIVED AT LEAST TWO PRIOR SYSTEMIC THERAPIES
- ODE-72 TREATMENT OF PATIENTS WITH CHRONIC LYMPHOCYTIC LEUKEMIA WITH 17P DELETION WHO HAVE NOT RECEIVED AT LEAST ONE PRIOR THERAPY
- ODE-73 LONG-TERM TREATMENT OF ADULT PATIENTS WITH GAUCHER DISEASE TYPE 1 WHO ARE CYP2D6 EXTENSIVE METABOLIZERS (EMS), INTERMEDIATE METABOLIZERS (IMS), OR POOR METABOLIZERS (PMS) AS DETECTED BY AN FDA-CLEARED TEST.
- ODE-74 TREATMENT OF PATIENTS WITH SEVERE APLASTIC ANEMIA WHO HAVE HAD AN INSUFFICIENT RESPONSE TO IMMUNOSUPPRESSIVE THERAPY
- ODE-75 TREATMENT OF PATIENTS WITH SEVERE APLASTIC ANEMIA WHO HAVE HAD AN INSUFFICIENT RESPONSE TO IMMUNOSUPPRESSIVE THERAPY.
- ODE-76 TREATMENT OF PATIENTS WITH MANTLE CELL LYMPHOMA WHO HAVE NOT RECEIVED AT LEAST 1 PRIOR THERAPY
- ODE-77 TREATMENT OF IDIOPATHIC PULMONARY FIBROSIS
- ODE-78 TREATMENT OF HYPERCALCEMIA IN ADULT PATIENTS WITH PRIMARY HYPERPARATHYROIDISM FOR WHOM PARATHYROIDECTOMY WOULD BE INDICATED ON THE BASIS OF SERUM CALCIUM LEVELS, BUT WHO ARE UNABLE TO UNDERGO PARATHYROIDECTOMY.
- ODE-79 TREATMENT OF PATIENTS WITH POLYCYTHEMIA VERA WHO HAVE HAD AN INADEQUATE RESPONSE TO OR ARE INTOLERANT OF HYDROXYUREA
- ODE-80 TREATMENT OF PEDIATRIC PATIENTS WITH TOURETTE'S
- ODE-81 TREATMENT OF PATIENTS WITH ACROMEGALY WHO HAVE HAD AN INADEQUATE RESPONSE TO SURGERY AND/OR FOR WHOM SURGERY IS NOT AN OPTION
- ODE-82 TREATMENT OF PATIENTS WITH UNRESECTABLE, WELL- OR MODERATELY-DIFFERENTIATED LOCALLY ADVANCED OR METASTATIC GASTROENTEROPANCREATIC NEUROENDOCRINE TUMORS TO IMPROVE PROGRESSION-FREE SURVIVAL
- ODE-83 USE OF AS MONOTHERAPY FOR PATIENTS WITH DELETERIOUS OR SUSPECTED DELETERIOUS GERMLINE BRCA MUTATED (AS DETECTED BY AN FDA-APPROVED TEST) ADVANCED OVARIAN CANCER WHO HAVE BEEN TREATED WITH THREE OR MORE PRIOR LINES OF CHEMOTHERAPY
- ODE-84 TREATMENT OF MOTOR FLUCTUATIONS IN PATIENTS WITH ADVANCED PARKINSON'S DISEASE
- ODE-85 AS A REPLACEMENT SOLUTION IN CONTINUOUS RENAL REPLACEMENT THERAPY (CRRT) AND IN CASE OF DRUG POISONING WHEN CRRT IS USED TO REMOVE DIALYZABLE SUBSTANCES
- ODE-86 TREATMENT OF PATIENTS WITH WALDENSTROM'S MACROGLOBULINEMIA
- ODE-87 TREATMENT OF PATIENTS WITH LOCALLY RECURRENT OR METASTATIC, PROGRESSIVE, RADIOACTIVE IODINE REFRACTORY DIFFERENTIATED THYROID CANCER
- ODE-88 FOR USE IN COMBINATION WITH DEXAMETHASONE FOR THE TREATMENT OF PATIENTS WITH MULTIPLE MYELOMA WHO HAVE NOT RECEIVED AT LEAST ONE PRIOR THERAPY (FIRST LINE TREATMENT)
- ODE-89 TREATMENT OF PATIENTS WITH MULTIPLE MYELOMA WHO HAVE RECEIVED AT LEAST 2 PRIOR REGIMENS, INCLUDING BORTEZOMIB AND AN IMMUNOMODULATORY AGENT
- ODE-90 TREATMENT OF INVASIVE MUCORMYCOSIS IN PATIENTS 18 YEARS OF AGE AND OLDER
- ODE-91 TREATMENT OF BILE ACID SYNTHESIS DISORDERS DUE TO SINGLE ENZYME DEFECTS
- ODE-92 TREATMENT OF LYMPHANGIOLEIOMYOMATOSIS (LAM)
- ODE-93 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGE 12 YEARS AND OLDER WHO ARE HOMOZYGOUS FOR F508DEL MUTATION IN THE CFTR GENE
- ODE-94 PROPHYLAXIS OF ORGAN REJECTION IN KIDNEY TRANSPLANT PATIENTS CONVERTED FROM TACROLIMUS IMMEDIATE-RELEASE FORMULATIONS IN COMBINATION WITH OTHER IMMUNOSUPPRESSANTS
- ODE-95 FOR THE FIRST-LINE TREATMENT OF PATIENTS WITH METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) WHOSE TUMORS HAVE EPIDERMAL GROWTH FACTOR RECEPTOR (EGFR) EXON 19 DELETIONS OR EXON 21 (L858R) SUBSTITUTION MUTATIONS AS DETECTED BY AN FDA-APPROVED TEST

PATENT AND EXCLUSIVITY TERMS**ORPHAN DRUG EXCLUSIVITY**

- ODE-96 TREATMENT OF PRIMARY HYPERKALEMIC PERIODIC PARALYSIS, PRIMARY HYPOKALEMIC PERIOD PARALYSIS, AND RELATED VARIANTS
- ODE-97 TO EXPAND THE INDICATION TO PEDIATRIC PATIENTS 2-6 YEARS OF AGE WITH NEPHROPATHIC CYSTINOSIS
- ODE-98 TREATMENT OF HEREDITARY OROTIC ACIDURIA
- ODE-99 FOR USE IN COMBINATION WITH 5-FLUOROURACIL AND LEUCOVORIN, FOR THE TREATMENT OF PATIENTS WITH METASTATIC ADENOCARCINOMA OF THE PANCREAS THAT HAS PROGRESSED FOLLOWING GEMCITABINE-BASED THERAPY
- ODE-100 FOR THE TREATMENT OF PATIENTS WITH UNRESECTABLE OR METASTATIC LIPOSARCOMA OR LEIOMYOSARCOMA WHO RECEIVED A PRIOR ANTHRACYCLINE-CONTAINING REGIMEN
- ODE-101 FOR THE TREATMENT OF PATIENTS WITH UNRESECTABLE OR METASTATIC MELANOMA WITH BRAF V600E OR V600K MUTATION, IN COMBINATION WITH VEMURAFENIB. COTELLIC IS NOT INDICATED FOR TREATMENT OF PATIENTS WITH WILD-TYPE BRAF MELANOMA
- ODE-102 FOR TREATMENT OF PATIENTS WITH METASTATIC EPIDERMAL GROWTH FACTOR RECEPTOR (EGFR) T790M MUTATION-POSITIVE NON-SMALL CELL LUNG CANCER (NSCLC), AS DETECTED BY AN FDA-APPROVED TEST, WHO HAVE PROGRESSED ON OR AFTER EGFR TKI THERAPY
- ODE-103 USE IN COMBINATION WITH LENALIDOMIDE AND DEXAMETHASONE FOR THE TREATMENT OF PATIENTS WITH MULTIPLE MYELOMA WHO HAVE RECEIVED AT LEAST ONE PRIOR THERAPY
- ODE-104 EMERGENCY TX OF PTS FOLLOWING A FU OR CAPECITABINE OD, OR WHO EXHIBIT EARLY-ONSET, SEVERE OR LIFE-THREATENING TOXICITY AFFECTING THE CARDIAC SYSTEM OR CNS, AND/OR EARLY-ONSET, UNUSUALLY SEVERE AR W/IN 96 HRS FOLLOWING THE END OF FU OR CAPECITABINE ADMIN.
- ODE-105 TREATMENT OF PATIENTS WITH ANAPLASTIC LYMPHOMA KINASE (ALK)-POSITIVE METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC), WHO HAVE PROGRESSED ON OR ARE INTOLERANT TO CRIZOTINIB
- ODE-106 FOR USE OF UPTRAVI (SELEXIPAG) TABLETS, 200, 400, 600, 800, 1000, 1200, 1400, AND 1600 MCG FOR TREATMENT OF PULMONARY ARTERIAL HYPERTENSION (PAH, WHO GROUP I) TO REDUCE THE RISKS OF DISEASE PROGRESSION AND HOSPITALIZATION FOR PAH
- ODE-107 TREATMENT OF PATIENTS WITH UNRESECTABLE OR METASTATIC LIPOSARCOMA WHO HAVE RECEIVED A PRIOR ANTHRACYCLINE-CONTAINING REGIMEN
- ODE-108 TREATMENT OF ADULT PATIENTS WITH PROGRESSIVE, WELL-DIFFERENTIATED, NON-FUNCTIONAL, NEUROENDOCRINE TUMORS (NET) OF GASTROINTESTINAL (GI) OR LUNG ORIGIN, (EXCLUDING PANCREATIC) WITH UNRESECTABLE, LOCALLY ADVANCED OR METASTATIC DISEASE
- ODE-109 INDICATED FOR THE TREATMENT OF PATIENTS WITH CHRONIC LYMPHOCYTIC LEUKEMIA WITHOUT 17P DELETION WHO HAVE NOT RECEIVED AT LEAST ONE PRIOR THERAPY (FIRST LINE THERAPY)
- ODE-110 FOR HIGH-DOSE CONDITIONING TREATMENT PRIOR TO HEMATOPOIETIC PROGENITOR (STEM) CELL TRANSPLANTATION IN PATIENTS WITH MULTIPLE MYELOMA
- ODE-111 TREATMENT OF PATIENTS WITH METASTATIC NON-SMALL CELL LUNG CANCER WHOSE TUMORS ARE ROS-1 POSITIVE.
- ODE-112 FOR THE TREATMENT OF ADULT AND PEDIATRIC PATIENTS WITH HEPATIC VENO-OCCLUSIVE DISEASE (VOD), ALSO KNOWN AS SINUSOIDAL OBSTRUCTION SYNDROME (SOS), WITH RENAL OR PULMONARY DYSFUNCTION FOLLOWING HEMATOPOIETIC STEM CELL TRANSPLANTATION (HSCT).
- ODE-113 FOR TREATMENT OF PEDIATRIC AND ADULT PATIENTS WITH ACQUIRED METHEMOGLOBINEMIA.
- ODE-114 TREATMENT OF PATIENTS WITH CHRONIC LYMPHOCYTIC LEUKEMIA (CLL) WITH 17P DELETION, AS DETECTED BY AN FDA APPROVED TEST, WHO HAVE RECEIVED AT LEAST ONE PRIOR THERAPY
- ODE-115 TREATMENT OF PATIENTS WITH METASTATIC, SQUAMOUS, NON-SMALL CELL LUNG CANCER PROGRESSED AFTER PLATINUM-BASED CHEMOTHERAPY
- ODE-116 TREATMENT OF PROGRESSIVE KERATOCONUS
- ODE-117 FOR TREATMENT OF PATIENTS WITH SMALL LYMPHOCYTIC LYMPHOMA (SLL)
- ODE-118 AN ADJUNCT TO DIET TO REDUCE LDL-C, TOTAL-C, NONHDL-C AND APOB IN CHILDREN AND ADOLESCENTS 7 TO 17 YEARS OF AGE WITH HOMOZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA, EITHER ALONE OR WITH OTHER LIPID-LOWERING TREATMENTS (E.G., LDL APHERESIS)
- ODE-119 TREATMENT OF PRIMARY BILIARY CHOLANGITIS (PBC) IN COMBINATION WITH URSODEOXYCHOLIC ACID (UDCA) IN ADULTS WITH AN INADEQUATE RESPONSE TO UDCA, OR AS MONOTHERAPY IN ADULTS UNABLE TO TOLERATE UDCA

PATENT AND EXCLUSIVITY TERMS**ORPHAN DRUG EXCLUSIVITY**

ODE-120	FOR USE AFTER RADIOLABELING WITH GA 68, WITH POSITRON EMISSION TOMOGRAPHY (PET) FOR LOCALIZATION OF SOMATOSTATIN RECEPTOR POSITIVE NEUROENDOCRINE TUMORS (NETS) IN ADULT AND PEDIATRIC PATIENTS.
ODE-121	TREATMENT OF CORNEAL ECTASIA FOLLOWING REFRACTIVE SURGERY
ODE-122	TREATMENT OF DUCHENNE MUSCULAR DYSTROPHY (DMD) IN PATIENTS WHO HAVE A CONFIRMED MUTATION OF THE DMD GENE THAT IS AMENABLE TO EXON 51 SKIPPING
ODE-123	TREATMENT OF CYSTIC FIBROSIS (CF) IN PATIENTS AGE 6-11 YEAR OLD WHO ARE HOMOZYGOUS FOR THE F508DEL MUTATION IN THE CFTR GENE
ODE-124	REPLACEMENT THERAPY FOR ORAL CARBAMAZEPINE FORMULATIONS, WHEN ORAL ADMINISTRATION IS TEMPORARILY NOT FEASIBLE, IN ADULTS WITH THE FOLLOWING SEIZURE TYPES: PARTIAL WITH COMPLEX SYMPTOMOLOGY, GENERALIZED CLONIC-TONIC, AND MIXED
ODE-125	INDICATED IN PEDIATRIC PATIENTS 10 YEARS AND OLDER FOR THE PREVENTION AND TREATMENT OF SECONDARY HYPERPARATHYROIDISM ASSOCIATED WITH CHRONIC KIDNEY DISEASE (CKD) STAGES 3 AND 4 AND CKD STAGE 5 IN PATIENTS ON HEMODIALYSIS OR PERITONEAL DIALYSIS
ODE-126	AS MONOTHERAPY FOR THE TREATMENT OF PATIENTS WITH DELETERIOUS BRCA MUTATION (GERMLINE AND/OR SOMATIC) ASSOCIATED ADVANCED OVARIAN CANCER WHO HAVE BEEN TREATED WITH TWO OR MORE CHEMOTHERAPIES
ODE-127	TREATMENT OF SPINAL MUSCULAR ATROPHY IN PEDIATRIC AND ADULT PATIENTS
ODE-128	TREATMENT OF PATIENTS WITH MARGINAL ZONE LYMPHOMA (MZL) WHO REQUIRE SYSTEMIC THERAPY AND HAVE RECEIVED AT LEAST ONE PRIOR ANTI-CD20-BASED THERAPY
ODE-129	INDICATED FOR REDUCING THE RISK OF GRAFT REJECTION WHEN USED IN CONJUNCTION WITH HIGH-DOSE BUSULFAN & CYCLOPHOSPHAMIDE AS A PREPARATIVE REGIMEN FOR ALLOGENIC HEMATOPOIETIC PROGENITOR CELL TRANSPLANTATION FOR PEDIATRIC PATIENTS WITH CLASS 3 BETA-THALASSEMIA
ODE-130	TREATMENT OF DUCHENNE MUSCULAR DYSTROPHY IN PATIENTS 5 YEARS OF AGE AND OLDER
ODE-131	TREATMENT OF MULTIPLE MYELOMA (MM), AS MAINTENANCE FOLLOWING AUTOLOGOUS HEMATOPOIETIC STEM CELL TRANSPLANTATION (AUTO-HSCT)
ODE-132	TREATMENT OF CARCINOID SYNDROME DIARRHEA IN COMBINATION WITH SOMATOSTATIN ANALOG (SSA) THERAPY IN ADULTS INADEQUATELY CONTROLLED BY SSA THERAPY
ODE-133	INDICATED FOR MAINTENANCE TREATMENT OF ADULT PATIENTS WITH RECURRENT EPITHELIAL OVARIAN, FALLOPIAN TUBE, OR PRIMARY PERITONEAL CANCER WHO ARE IN A COMPLETE OR PARTIAL RESPONSE TO PLATINUM-BASED CHEMOTHERAPY
ODE-134	TREATMENT OF CHOREA ASSOCIATED WITH HUNTINGTON'S DISEASE
ODE-135	TREATMENT OF CHRONIC HCV GENOTYPE 2 OR 3 INFECTION IN PEDIATRIC PATIENTS 12 YEARS OF AGE AND OLDER OR WEIGHING AT LEAST 35 KG WITHOUT CIRRHOSIS OR WITH COMPENSATED CIRRHOSIS FOR USE IN COMBINATION WITH RIBAVIRIN
ODE-136	TREATMENT OF PEDIATRIC PATIENTS 12 YEARS OF AGE AND OLDER OR WEIGHING AT LEAST 35 KG WITH CHRONIC HEPATITIS C VIRUS GENOTYPE 1, 4, 5, OR 6 INFECTION WITHOUT CIRRHOSIS OR WITH COMPENSATED CIRRHOSIS
ODE-137	TREATMENT OF OLIGOARTICULAR JUVENILE IDIOPATHIC ARTHRITIS (PERSISTENT OLIGOARTHRITIS, PSORIATIC JUVENILE IDIOPATHIC ARTHRITIS, ENTHESITIS-RELATED ARTHRITIS, OR UNDIFFERENTIATED ARTHRITIS) & POLYARTICULAR JUVENILE IDIOPATHIC ARTHRITIS IN CHILDREN 0-16 YRS
ODE-138	TREATMENT OF PEDIATRIC PATIENTS WITH ACUTE LYMPHOBLASTIC LEUKEMIA AS A COMPONENT OF A COMBINATION CHEMOTHERAPY MAINTENANCE REGIMEN
ODE-139	TREATMENT OF PATIENTS WITH HEPATOCELLULAR CARCINOMA (HCC OR LIVER CANCER) WHO HAVE BEEN PREVIOUSLY TREATED WITH THE DRUG SORAFENIB.
ODE-140	TREATMENT OF ADULT PATIENTS WITH AGGRESSIVE SYSTEMIC MASTOCYTOSIS (ASM), SYSTEMIC MASTOCYTOSIS WITH ASSOCIATED HEMATOLOGICAL NEOPLASM (SM-AHN), OR MAST CELL LEUKEMIA (MCL)
ODE-141	TREATMENT OF ADULT PATIENTS WITH NEWLY DIAGNOSED ACUTE MYELOID LEUKEMIA (AML) THAT IS FLT3 MUTATION-POSITIVE AS DETECTED BY AN FDA APPROVED TEST, IN COMBINATION WITH STANDARD CYTARABINE AND DAUNORUBICIN INDUCTION AND CYTARABINE CONSOLIDATION
ODE-142	TREATMENT OF PATIENTS WITH ANAPLASTIC LYMPHOMA KINASE (ALK)-POSITIVE METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) WHO HAVE PROGRESSED ON OR ARE INTOLERANT TO CRIZOTINIB
ODE-143	TO DECREASE THE RECURRENCE OF PNEUMOTHORAX IN ADULTS

PATENT AND EXCLUSIVITY TERMS**ORPHAN DRUG EXCLUSIVITY**

- ODE-144 TREATMENT OF AMYOTROPHIC LATERAL SCLEROSIS (ALS)
- ODE-145 TREATMENT OF PATIENTS WITH METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) WHOSE TUMORS ARE ANAPLASTIC LYMPHOMA KINASE (ALK)-POSITIVE AS DETECTED BY AN FDA-APPROVED TEST
- ODE-146 OPTICAL IMAGING AGENT INDICATED IN PATIENTS WITH GLIOMA (SUSPECTED WORLD HEALTH ORGANIZATION GRADES III OR IV ON PREOPERATIVE IMAGING) AS AN ADJUNCT FOR THE VISUALIZATION OF MALIGNANT TISSUE DURING SURGERY
- ODE-147 DABRAFENIB IN COMBINATION WITH TRAMETINIB, FOR THE TX. OF PTS WITH METASTATIC NON-SMALL CELL LUNG CANCER WITH BRAF V600E MUTATION AS DETECTED BY AN FDA-APPROVED TEST
- ODE-148 TRAMETINIB IN COMBINATION WITH DABRAFENIB, FOR THE TX. OF PTS WITH METASTATIC NON-SMALL CELL LUNG CANCER WITH BRAF V600E MUTATION AS DETECTED BY AN FDA-APPROVED TEST
- ODE-149 TREATMENT OF PEDIATRIC PATIENTS 2 YEARS OF AGE AND OLDER WITH CENTRAL PRECOCIOUS PUBERTY
- ODE-150 TO REDUCE THE ACUTE COMPLICATIONS OF SICKLE CELL DISEASE IN ADULT AND PEDIATRIC PATIENTS 5 YEARS OF AGE AND OLDER.
- ODE-151 TREATMENT OF ADULT PATIENTS WITH RELAPSED OR REFRACTORY ACUTE MYELOID LEUKEMIA WITH AN ISOCITRATE DEHYDROGENASE-2 (IDH2) MUTATION AS DETECTED BY AN FDA-APPROVED TEST
- ODE-152 TREATMENT OF ADULT PATIENTS WITH CHRONIC GRAFT VERSUS HOST DISEASE (CGVHD)
- ODE-153 TREATMENT OF DYSKINESIA IN PATIENTS WITH PARKINSON'S DISEASE RECEIVING LEVODOPA-BASED THERAPY WITH OR WITHOUT CONCOMITANT DOPAMINERGIC MEDICATIONS
- ODE-154 FOR USE IN CHILDREN AGES 2 TO 12 YEARS OLD WITH CHAGAS DISEASE
- ODE-155 TREATMENT OF ADULT PATIENTS WITH RELAPSED FOLLICULAR LYMPHOMA WHO HAVE RECEIVED AT LEAST TWO PRIOR SYSTEMIC THERAPIES
- ODE-156 TREATMENT OF ADULTS WITH CARCINOID SYNDROME; WHEN USED, IT REDUCES THE FREQUENCY OF SHORT-ACTING SOMATOSTATIN ANALOG RESCUE THERAPY
- ODE-157 FOR USE AS A NITROGEN-BINDING AGENT FOR CHRONIC MANAGEMENT OF PEDIATRIC PATIENTS ≥ 2 MONTHS AND < 2 YEARS OF AGE WITH UREA CYCLE DISORDERS (UCDS) WHO CANNOT BE MANAGED BY DIETARY PROTEIN RESTRICTION AND/OR AMINO ACID SUPPLEMENTATION ALONE
- ODE-158 TREATMENT OF PATIENTS WITH ERDHEIM-CHESTER DISEASE WITH BRAF V600 MUTATION
- ODE-159 FOR TREATMENT OF PATIENTS WITH ANAPLASTIC LYMPHOMA KINASE (ALK) POSITIVE, METASTATIC NON-SMALL-CELL LUNG CANCER (NSCLC) AS DETECTED BY AN FDA APPROVED TEST, EXCLUDING PATIENTS WHO HAVE PROGRESSED ON OR ARE INTOLERANT TO CRIZOTINIB
- ODE-160 FOR TREATMENT OF SCURVY IN ADULT AND PEDIATRIC PATIENTS AGE 5 MONTHS AND OLDER FOR WHOM ORAL ADMINISTRATION IS NOT POSSIBLE, INSUFFICIENT OR CONTRAINDICATED
- ODE-161 TREATMENT OF PULMONARY ARTERIAL HYPERTENSION (PAH) (WHO GROUP 1) IN PEDIATRIC PATIENTS AGED 3 YRS AND OLDER WITH IDIOPATHIC OR CONGENITAL PAH TO IMPROVE PULMONARY VASCULAR RESISTANCE (PVR), WHICH IS EXPECTED TO RESULT IN AN IMPROVEMENT IN EXERCISE ABILITY
- ODE-162 TREATMENT OF NEPHROPATHIC CYSTINOSIS IN PEDIATRIC PATIENTS 1 YEAR OF AGE TO LESS THAN 2 YEARS OF AGE
- ODE-163 TREATMENT OF ADULT PATIENTS WITH NEWLY-DIAGNOSED CHRONIC PHASE (CP) PHILADELPHIA CHROMOSOME-POSITIVE CHRONIC MYELOGENOUS LEUKEMIA (PH+ CML)
- ODE-164 TREATMENT OF PEDIATRIC PATIENTS WITH PHILADELPHIA CHROMOSOME-POSITIVE (PH+) CHRONIC MYELOID LEUKEMIA (CML) IN CHRONIC PHASE
- ODE-165 PROPHYLAXIS OF CYTOMEGALOVIRUS (CMV) INFECTION AND DISEASE IN ADULT CMV-SEROPOSITIVE RECIPIENTS [R+] OF AN ALLOGENEIC HEMATOPOIETIC STEM CELL TRANSPLANT (HSCT)
- ODE-166 TREATMENT OF SOMATOSTATIN RECEPTOR-POSITIVE GASTROENTEROPANCREATIC NEUROENDOCRINE TUMORS (GEP-NETS) INCLUDING FOREGUT, MIDGUT, AND HINDGUT NEUROENDOCRINE TUMORS IN ADULTS
- ODE-167 ARSENIC TRIOXIDE FOR USE IN COMBINATION WITH TRETINOIN FOR TREATMENT OF ADULTS WITH NEWLY-DIAGNOSED LOW-RISK ACUTE PROMYELOCYTIC LEUKEMIA (APL) WHOSE APL IS CHARACTERIZED BY THE PRESENCE OF THE T(15;17) TRANSLOCATION OR PML/RAR-ALPHA GENE EXPRESSION
- ODE-168 FOR THE MAINTENANCE TREATMENT OF ADULT PATIENTS WITH RECURRENT EPITHELIAL OVARIAN, FALLOPIAN TUBE, OR PRIMARY PERITONEAL CANCER WHO ARE IN A COMPLETE OR

PATENT AND EXCLUSIVITY TERMS**ORPHAN DRUG EXCLUSIVITY**

PARTIAL RESPONSE TO PLATINUM-BASED CHEMOTHERAPY

- ODE-169 FOR THE ADJUNCTIVE TREATMENT OF ADULT AND PEDIATRIC PATIENTS AGED 2 YEARS AND OLDER WITH TUBEROUS SCLEROSIS COMPLEX (TSC)-ASSOCIATED PARTIAL-ONSET SEIZURES
- ODE-170 FOR THE DIAGNOSIS OF ADULT GROWTH HORMONE DEFICIENCY (AGHD)
- ODE-171 TREATMENT OF PEDIATRIC PATIENTS GREATER THAN OR EQUAL TO 1 YEAR OF AGE WITH NEWLY DIAGNOSED PHILADELPHIA CHROMOSOME POSITIVE CHRONIC MYELOID LEUKEMIA (PH+CML) IN CHRONIC PHASE
- ODE-172 TREATMENT OF PEDIATRIC PATIENTS GREATER THAN OR EQUAL TO 1 YEAR OF AGE WITH CHRONIC PHASE PHILADELPHIA CHROMOSOME POSITIVE CHRONIC MYELOID LEUKEMIA WITH RESISTANCE OR INTOLERANCE TO PRIOR TYROSINE-KINASE INHIBITOR THERAPY
- ODE-173 TREATMENT OF PATIENTS WITH CYSTIC FIBROSIS AGED 12 YEARS AND OLDER WHO ARE HOMOZYGOUS FOR THE F508DEL MUTATION OR WHO HAVE AT LEAST ONE MUTATION IN THE CYSTIC FIBROSIS TRANSMEMBRANE CONDUCTANCE REGULATOR GENE RESPONSIVE TO TEZACAFTOR/IVACAFTOR
- ODE-174 FOR THE TREATMENT OF THROMBOCYTOPENIA IN ADULT PATIENTS WITH CHRONIC IMMUNE THROMBOCYTOPENIA (ITP) WHO HAVE HAD AN INSUFFICIENT RESPONSE TO A PREVIOUS TREATMENT
- ODE-175 TREATMENT OF ADULT PATIENTS WITH MANTLE CELL LYMPHOMA (MCL) WHO HAVE RECEIVED AT LEAST ONE PRIOR THERAPY
- ODE-176 INDICATED FOR THE FIRST-LINE TREATMENT OF PATIENTS WITH METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) WHOSE TUMORS HAVE EPIDERMAL GROWTH FACTOR RECEPTOR (EGFR) EXON 19 DELETIONS OR EXON 21 L858R MUTATIONS, AS DETECTED BY AN FDA-APPROVED TEST
- ODE-177 TO REDUCE THE FREQUENCY OF PAINFUL CRISES AND TO REDUCE THE NEED FOR BLOOD TRANSFUSIONS IN PEDIATRIC PATIENTS, 2 YEARS OF AGE AND OLDER, WITH SICKLE CELL ANEMIA WITH RECURRENT MODERATE TO SEVERE PAINFUL CRISIS
- ODE-178 INDICATED TO SLOW KIDNEY FUNCTION DECLINE IN ADULTS AT RISK OF RAPIDLY PROGRESSING AUTOSOMAL DOMINANT POLYCYSTIC KIDNEY DISEASE (ADPKD)
- ODE-179 TREATMENT OF PATIENTS WITH CLL AND TREATMENT OF PATIENTS WITH INDOLENT B-CELL NHL THAT HAS PROGRESSED DURING OR WITHIN SIX MONTHS OF TREATMENT WITH RITUXIMAB OR A RITUXIMAB CONTAINING REGIMEN
- ODE-180 MAINTENANCE TREATMENT OF ADULT PATIENTS WITH RECURRENT EPITHELIAL OVARIAN, FALLOPIAN TUBE OR PRIMARY PERITONEAL CANCER, WHO ARE IN A COMPLETE OR PARTIAL RESPONSE TO PLATINUM-BASED CHEMOTHERAPY
- ODE-181 TREATMENT OF ADULT PATIENTS WITH DELETERIOUS OR SUSPECTED DELETERIOUS GERMLINE BRCA-MUTATED ADVANCED OVARIAN CANCER WHO HAVE BEEN TREATED WITH THREE OR MORE PRIOR LINES OF CHEMOTHERAPY
- ODE-182 TRAMETINIB IS INDICATED, IN COMBINATION WITH DABRAFENIB, FOR THE ADJUVANT TREATMENT OF PATIENTS WITH MELANOMA WITH BRAF V600E OR V600K MUTATIONS, AS DETECTED BY AN FDA-APPROVED TEST, AND INVOLVEMENT OF LYMPH NODE(S), FOLLOWING COMPLETE RESECTION
- ODE-183 TRAMETINIB AND DABRAFENIB IN COMBINATION, FOR THE TREATMENT OF PATIENTS WITH LOCALLY ADVANCED OR METASTATIC ANAPLASTIC THYROID CANCER (ATC) WITH BRAF V600E MUTATION AND WITH NO SATISFACTORY LOCOREGIONAL TREATMENT OPTIONS
- ODE-184 INDICATED IN COMBINATION WITH OTHER ANTIRETROVIRAL AGENTS FOR THE TREATMENT OF PEDIATRIC PATIENTS WITH HIV-1 INFECTION
- ODE-185 INDICATED FOR THE TREATMENT OF PATIENTS WITH CHRONIC LYMPHOCYTIC LEUKEMIA (CLL) OR SMALL LYMPHOCYTIC LYMPHOMA (SLL), WITH OR WITHOUT 17P DELETION, WHO HAVE RECEIVED AT LEAST ONE PRIOR THERAPY
- ODE-186 TREATMENT OF CYSTIC FIBROSIS (CF) IN PATIENTS AGE 6 YEARS AND OLDER WHO HAVE ONE OF THE FOLLOWING MUTATIONS IN THE CFTR GENE: G1244E, G1349D, G178R, G551S, S1251N, S1255P, S549N, OR S549R
- ODE-187 TREATMENT OF CYSTIC FIBROSIS (CF) IN PATIENTS AGE 6 YEARS AND OLDER WHO HAVE AN R117H MUTATION IN THE CFTR GENE
- ODE-188 TREATMENT OF CYSTIC FIBROSIS (CF) IN PATIENTS AGES 2 TO LESS THAN 6 YEARS WHO HAVE ONE OF THE FOLLOWING MUTATIONS IN THE CFTR GENE: G551D, G1244E, G1349D, G178R, G551S, S1251N, S1255P, S549N, S549R, AND R117H
- ODE-189 TREATMENT OF CYSTIC FIBROSIS (CF) IN PATIENTS AGE 2 YEARS AND OLDER WHO HAVE ONE OF THE FOLLOWING MUTATIONS IN THE CFTR GENE: 711+3A-G, E831X, 2789+5G-A, 3272-26A-G, AND 3849+10KBC-T

PATENT AND EXCLUSIVITY TERMS**ORPHAN DRUG EXCLUSIVITY**

- ODE-190 TX OF CF IN PTS 2 YRS AND OLDER WHO HAVE ONE OF THE FOLLOWING MUTATIONS IN THE CFTR GENE: E56K, P67L, R74W, D110E, D110H, R117C, E193K, L206W, R347H, R352Q, A455E, D579G, S945L, S977F, F1052V, K1060T, A1067T, G1069R, R1070Q, R1070W, F1074L, D1152H, D1270N
- ODE-191 TO DECREASE THE RECURRENCE OF MALIGNANT PLEURAL EFFUSIONS IN SYMPTOMATIC PATIENTS FOLLOWING MAXIMAL DRAINAGE OF THE PLEURAL EFFUSION
- ODE-192 INDICATED TO INDUCE CONTROLLED CARDIAC SEPTAL INFRACTION TO IMPROVE EXERCISE CAPACITY IN ADULTS WITH SYMPTOMATIC HYPERTROPHIC OBSTRUCTIVE CARDIOMYOPATHY WHO ARE NOT CANDIDATES FOR SURGICAL MYECTOMY
- ODE-193 INDICATED FOR THE TREATMENT OF ONCHOCERCIASIS DUE TO ONCHOCERCA VOLVULUS IN PATIENTS AGED 12 YEARS AND OLDER
- ODE-194 ENCORAFENIB IS INDICATED IN COMBINATION WITH BINIMETINIB, FOR THE TREATMENT OF PATIENTS WITH UNRESECTABLE OR METASTATIC MELANOMA WITH A BRAF V600E OR V600K MUTATION, AS DETECTED BY AN FDA-APPROVED TEST
- ODE-195 FOR THE TREATMENT OF CYSTIC FIBROSIS (CF) IN PATIENTS AGE 2 THROUGH 5 YEARS OLD WHO ARE HOMOZYGOUS FOR THE F508DEL MUTATION IN THE CFTR GENE
- ODE-196 INDICATED FOR THE FIRST-LINE TREATMENT OF PATIENTS WITH UNRESECTABLE HEPATOCELLULAR CARCINOMA (HCC)
- ODE-197 INDICATED FOR THE TREATMENT OF THE POLYNEUROPATHY OF HEREDITARY TRANSTHYRETIN-MEDIATED AMYLOIDOSIS IN ADULTS
- ODE-198 INDICATED FOR THE TREATMENT OF SEIZURES ASSOCIATED WITH DRAVET SYNDROME (DS) IN PATIENTS 2 YEARS OF AGE AND OLDER TAKING CLOBAZAM
- ODE-199 THE TREATMENT OF CYSTIC FIBROSIS (CF) IN PATIENTS AGE 12 MONTHS AND OLDER WHO HAVE ONE MUTATION IN THE CFTR GENE THAT IS RESPONSIVE TO IVACAFTOR POTENTIATION BASED ON CLINICAL AND/OR IN VITRO ASSAY DATA
- ODE-200 INDICATED FOR THE TREATMENT OF HUMAN SMALLPOX DISEASE CAUSED BY VARIOLA VIRUS IN ADULTS AND PEDIATRIC PATIENTS WEIGHING AT LEAST 13 KG
- ODE-201 INDICATED FOR THE RADICAL CURE (PREVENTION OF RELAPSE) OF PLASMODIUM VIVAX MALARIA IN PATIENTS AGED 16 YEARS AND OLDER WHO ARE RECEIVING APPROPRIATE ANTIMALARIAL THERAPY FOR ACUTE P. VIVAX INFECTION
- ODE-202 INDICATED AS A SOURCE OF CALORIES AND FATTY ACIDS IN PEDIATRIC PATIENTS WITH PARENTERAL NUTRITION-ASSOCIATED CHOLESTASIS (PNAC)
- ODE-203 INDICATED FOR THE TREATMENT OF ADULT PATIENTS WITH RELAPSED OR REFRACTORY ACUTE MYELOID LEUKEMIA (AML) WITH A SUSCEPTIBLE ISOCITRATE DEHYDROGENASE-1 (IDH1) MUTATION AS DETECTED BY AN FDA-APPROVED TEST
- ODE-204 TREATMENT OF ADULT AND PEDIATRIC PATIENTS 12 YEARS AND OLDER WITH IOBENGUANE SCAN POSITIVE, UNRESECTABLE, LOCALLY ADVANCED OR METASTATIC PHEOCHROMOCYTOMA OR PARANGLIOMA WHO REQUIRE SYSTEMIC ANTICANCER THERAPY
- ODE-205 INDICATED FOR THE TREATMENT OF ADULTS WITH A CONFIRMED DIAGNOSIS OF FABRY DISEASE AND AN AMENABLE GALACTOSIDASE ALPHA GENE (GLA) VARIANT BASED ON IN VITRO ASSAY DATA
- ODE-206 FIRST-LINE TREATMENT OF PATIENTS WITH METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) WITH EPIDERMAL GROWTH FACTOR RECEPTOR (EGFR) EXON 19 DELETION OR EXON 21 L858R SUBSTITUTION MUTATIONS AS DETECTED BY AN FDA-APPROVED TEST
- ODE-207 TREATMENT OF STATUS EPILEPTICUS IN ADULTS
- ODE-208 TREATMENT OF ADULT PATIENTS WITH RELAPSED OR REFRACTORY CHRONIC LYMPHOCYTIC LEUKEMIA (CLL) OR SMALL LYMPHOCYTIC LYMPHOMA (SLL) AFTER AT LEAST TWO PRIOR THERAPIES
- ODE-209 TREATMENT OF ADULT PATIENTS WITH RELAPSED OR REFRACTORY FOLLICULAR LYMPHOMA (FL) AFTER AT LEAST TWO PRIOR SYSTEMIC THERAPIES
- ODE-210 INDICATED IN COMBINATION WITH STANDARD IMMUNOSUPPRESSIVE THERAPY FOR THE FIRST-LINE TREATMENT OF ADULT AND PEDIATRIC PATIENTS 2 YEARS AND OLDER WITH SEVERE APLASTIC ANEMIA
- ODE-211 INDICATED IN COMBO WITH AZACITIDINE, OR DECITABINE, OR LOW-DOSE CYTARABINE FOR THE TX OF NEWLY-DIAGNOSED ACUTE MYELOID LEUKEMIA IN ADULTS WHO ARE AGE 75 YEARS OR OLDER, OR WHO HAVE COMORBIDITIES THAT PRECLUDE USE OF INTENSIVE INDUCTION CHEMOTHERAPY
- ODE-212 INDICATED FOR THE TREATMENT OF THE POLYNEUROPATHY OF HEREDITARY TRANSTHYRETIN-MEDIATED AMYLOIDOSIS IN ADULTS

PATENT AND EXCLUSIVITY TERMS**ORPHAN DRUG EXCLUSIVITY**

- ODE-213 INDICATED FOR THE FIRST-LINE TREATMENT OF PATIENTS WITH METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) WITH EPIDERMAL GROWTH FACTOR (EGFR) EXON 19 DELETION OR EXON 21 L858R SUBSTITUTION MUTATIONS AS DETECTED BY AN FDA-APPROVED TEST
- ODE-214 TX OF MAC LUNG DISEASE IN ADULTS WITH LIMITED OR NO ALTERNATIVE TX OPTIONS AS PART OF A COMBO ANTIBACTERIAL DRUG REGIMEN WHO DO NOT ACHIEVE NEGATIVE SPUTUM CULTURES AFTER A MINIMUM OF 6 CONSECUTIVE MONTHS OF A MULTIDRUG BACKGROUND REGIMEN THERAPY
- ODE-215 INDICATED FOR THE TREATMENT OF ADULT AND PEDIATRIC PATIENTS WITH SOLID TUMORS THAT HAVE A NEUROTROPHIC RECEPTOR TYROSINE KINASE (NTRK) GENE FUSION WITHOUT A KNOWN ACQUIRED RESISTANCE MUTATION
- ODE-216 INDICATED FOR THE TREATMENT OF SEIZURES ASSOCIATED WITH LENNOX-GASTAUT SYNDROME (LGS) OR DRAVET SYNDROME (DS) IN PATIENTS 2 YEARS OF AGE AND OLDER
- ODE-217 INDICATED FOR THE TREATMENT OF PATIENTS WITH ANAPLASTIC LYMPHOMA KINASE (ALK)- POSITIVE METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) WHOSE DISEASE HAS PROGRESSED ON CRIZOTINIB AND AT LEAST ONE OTHER ALK INHIBITOR FOR METASTATIC DIESASE
- ODE-218 INDICATED FOR THE TREATMENT OF PATIENTS WITH ANAPLASTIC LYMPHOMA KINASE (ALK)- POSITIVE METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) WHOSE DISEASE HAS PROGRESSED ON ALECTINIB AS THE FIRST ALK INHIBITOR THERAPY FOR METASTATIC DISEASE
- ODE-219 INDICATED FOR THE TREATMENT OF PATIENTS WITH ANAPLASTIC LYMPHOMA KINASE (ALK)- POSITIVE METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) WHOSE DISEASE HAS PROGRESSED ON CERITINIB AS THE FIRST ALK INHIBITOR THERAPY FOR METASTATIC DISEASE
- ODE-220 INDICATED FOR THE TREATMENT OF ADULT AND PEDIATRIC PATIENTS WITH SOLID TUMORS THAT ARE METASTATIC OR WHERE SURGICAL RESECTION IS LIKELY TO RESULT IN SEVERE MORBIDITY
- ODE-221 INDICATED FOR THE TREATMENT OF ADULT AND PEDIATRIC PATIENTS WITH SOLID TUMORS THAT HAVE NO SATISFACTORY ALTERNATIVE TREATMENTS OR THAT HAVE PROGRESSED FOLLOWING TREATMENT
- ODE-222 INDICATED FOR THE TREATMENT OF ADULT PATIENTS WHO HAVE RELAPSED OR REFRACTORY ACUTE MYELOID LEUKEMIA (AML) WITH A FMS-LIKE TYROSINE KINASE 3 (FLT3) MUTATION AS DETECTED BY AN FDA-APPROVED TEST
- ODE-223 TREATMENT OF LAMBERT-EATON MYASTHENIC SYNDROME (LEMS)
- ODE-224 INDICATED, IN COMBINATION WITH LOW-DOSE CYTARABINE, FOR THE TREATMENT OF NEWLY-DIAGNOSED ACUTE MYELOID LEUKEMIA (AML) IN ADULT PATIENTS WHO ARE ≥ 75 YEARS OLD OR WHO HAVE COMORBIDITIES THAT PRECLUDE USE OF INTENSIVE INDUCTION CHEMOTHERAPY
- ODE-225 INDICATED FOR THE TREATMENT OF PEDIATRIC PATIENTS 1 YEAR OF AGE AND OLDER WITH NEWLY DIAGNOSED PHILADELPHIA CHROMOSOME-POSITIVE ACUTE LYMPHOBLASTIC LEUKEMIA (ALL) IN COMBINATION WITH CHEMOTHERAPY
- ODE-226 MAINTENANCE TREATMENT OF ADULTS WITH DELETERIOUS OR SUSPECTED DELETERIOUS GERMLINE OR SOMATIC BRCA-MUTATED ADVANCED EPITHELIAL OVARIAN, FALLOPIAN TUBE, OR PRIMARY PERITONEAL CANCER IN COMPLETE OR PARTIAL RESPONSE TO FIRST-LINE PLATINUM-BASED CHEMOTHERAPY
- ODE-227 INDICATED FOR TREATMENT OF PATIENTS WITH HEPATOCELLULAR CARCINOMA (HCC) WHO HAVE BEEN PREVIOUSLY TREATED WITH SORAFENIB
- ODE-228 INDICATED FOR THE TREATMENT OF FASCIOLIASIS IN PATIENTS 6 YEARS OF AGE AND OLDER
- ODE-229 TREATMENT OF ADULT PATIENTS WITH METASTATIC GASTRIC OR GASTROESOPHAGEAL JUNCTION ADENOCARCINOMA PREVIOUSLY TREATED WITH AT LEAST TWO PRIOR LINES OF CHEMOTHERAPY, AND IF APPROPRIATE, HER2/NEU-TARGETED THERAPY
- ODE-230 FIRST-LINE TREATMENT OF METASTATIC NON-SMALL CELL LUNG CANCER WHOSE TUMORS HAVE NON-RESISTANT EPIDERMAL GROWTH FACTOR (EGFR) MUTATIONS OTHER THAN EXON 19 DELETIONS OR EXON 21 (L858R) SUBSTITUTION MUTATIONS AS DETECTED BY AN FDA-APPROVED TEST
- ODE-231 INDICATED FOR THE TREATMENT OF CATAPLEXY OR EXCESSIVE DAYTIME SLEEPINESS (EDS) IN PEDIATRIC PATIENTS 7 YEARS OF AGE AND OLDER WITH NARCOLEPSY
- ODE-232 TREATMENT OF PEDIATRIC PATIENTS 12 YEARS AND OLDER OR WEIGHING AT LEAST 45 KG WITH CHRONIC HEPATITIS C VIRUS (HCV) GENOTYPE 1,2,3,4,5 OR 6 INFECTION WITHOUT CIRRHOSIS OR WITH COMPENSATED CIRRHOSIS (CHILD-PUGH A)
- ODE-233 TREATMENT OF PEDIATRIC PATIENTS 12 YEARS AND OLDER OR WEIGHING AT LEAST 45 KG

PATENT AND EXCLUSIVITY TERMS**ORPHAN DRUG EXCLUSIVITY**

WITH HCV GENOTYPE 1 INFECTION, WHO PREVIOUSLY HAVE BEEN TREATED WITH A REGIMEN CONTAINING AN HCV NS5A INHIBITOR OR AN NS3/4A PROTEASE INHIBITOR (PI), BUT NOT BOTH

- ODE-234 INDICATED FOR THE TREATMENT OF STABLE SYMPTOMATIC HEART FAILURE DUE TO DILATED CARDIOMYOPATHY (DCM) IN PEDIATRIC PATIENTS AGED 6 MONTHS AND OLDER, WHO ARE IN SINUS RHYTHM WITH AN ELEVATED HEART RATE
- ODE-235 INDICATED FOR THE TREATMENT OF ACUTE HERPETIC KERATITIS (DENDRITIC ULCERS) IN PATIENTS WITH HERPES SIMPLEX (HSV-1 AND HSV-2) VIRUS
- ODE-236 TREATMENT OF CYSTIC FIBROSIS (CF) IN PATIENTS AGE 6 MONTHS TO LESS THAN 12 MONTHS WHO HAVE ONE MUTATION IN THE CFTR GENE THAT IS RESPONSIVE TO IVACAFTOR POTENTIATION BASED ON CLINICAL AND/OR IN VITRO ASSAY DATA
- ODE-237 TREATMENT OF THE CARDIOMYOPATHY OF WILD TYPE OR HEREDITARY TRANSTHYRETIN-MEDIATED AMYLOIDOSIS (ATTR-CM) IN ADULTS TO REDUCE CARDIOVASCULAR MORTALITY AND CARDIOVASCULAR-RELATED HOSPITALIZATION
- ODE-238 TREATMENT OF STEROID-REFRACTORY ACUTE GRAFT-VERSUS-HOST DISEASE (GVHD) IN ADULT AND PEDIATRIC PATIENTS 12 YEARS AND OLDER
- ODE-239 TREATMENT OF PREVIOUSLY UNTREATED ADULT PATIENTS WITH CHRONIC LYMPHOCYTIC LEUKEMIA (CLL) OR SMALL LYMPHOCYTIC LYMPHOMA (SLL)
- ODE-240 TREATMENT OF PEDIATRIC PATIENTS 1 YEAR OF AGE AND OLDER WITH SHORT BOWEL SYNDROME (SBS) WHO ARE DEPENDENT ON PARENTERAL SUPPORT
- ODE-241 INDICATED IN COMBINATION WITH A RITUXIMAB PRODUCT FOR THE TREATMENT OF ADULT PATIENTS WITH PREVIOUSLY TREATED FOLLICULAR LYMPHOMA (FL)
- ODE-242 TX OF NEWLY-DIAGNOSED ACUTE MYELOID LEUKEMIA WITH A SUSCEPTIBLE ISOCITRATE DEHYDROGENASE-1 MUTATION AS DETECTED BY AN FDA-APPROVED TEST IN ADULT PTS WHO ARE >=75 YRS OLD OR WHO HAVE COMORBIDITIES THAT PRECLUDE USE OF INTENSIVE INDUCTION CHEMOTHERAPY
- ODE-243 ACUTE TREATMENT OF INTERMITTENT, STEREOTYPIC EPISODES OF FREQUENT SEIZURE ACTIVITY (I.E., SEIZURE CLUSTERS, ACUTE REPETITIVE SEIZURES) THAT ARE DISTINCT FROM A PATIENT'S USUAL SEIZURE PATTERN IN PATIENTS WITH EPILEPSY 12 YEARS OF AGE AND OLDER
- ODE-244 TREATMENT OF LAMBERT-EATON MYASTHENIC SYNDROME (LEMS) IN PATIENTS 6 TO LESS THAN 17 YEARS OF AGE
- ODE-245 INDICATED IN COMBINATION WITH A RITUXIMAB PRODUCT FOR THE TREATMENT OF ADULT PATIENTS WITH PREVIOUSLY TREATED MARGINAL ZONE LYMPHOMA (MZL)
- ODE-246 TREATMENT OF THROMBOCYTOPENIA IN ADULT PATIENTS WITH CHRONIC IMMUNE THROMBOCYTOPENIA WHO HAVE HAD AN INSUFFICIENT RESPONSE TO A PREVIOUS TREATMENT
- ODE-247 TX OF PTS W/ CYSTIC FIBROSIS (CF) AGE 6 TO <12 YRS WHO ARE HOMOZYGOUS FOR THE F508DEL MUTATION OR W/ AT LEAST 1 MUTATION IN CF TRANSMEMBRANE CONDUCTANCE REGULATORY GENE RESPONSIVE TO TEZACAFTOR/IVACAFTOR BASED ON IN VITRO DATA AND/OR CLINICAL EVIDENCE
- ODE-248 TREATMENT OF ADULT PATIENTS WITH ORAL ULCERS ASSOCIATED WITH BEHCETS DISEASE
- ODE-249 TREATMENT OF PATIENTS WITH CYSTIC FIBROSIS (CF) AGE 6 YEARS TO LESS THAN 12 YEARS WHO ARE HOMOZYGOUS FOR F508DEL MUTATION OR WHO HAVE AT LEAST ONE MUTATION IN THE CF TRANSMEMBRANE CONDUCTANCE REGULATOR GENE THAT IS RESPONSIVE TO TEZACAFTOR/IVACAFTOR
- ODE-250 INDICATED FOR THE TREATMENT OF ADULT PATIENTS WITH SYMPTOMATIC TENOSYNOVIAL GIANT CELL TUMOR (TGCT) ASSOCIATED WITH SEVERE MORBIDITY OR FUNCTIONAL LIMITATIONS AND NOT AMENABLE TO IMPROVEMENT WITH SURGERY
- ODE-251 INDICATED AS PART OF COMBINATION THERAPY IN THE TREATMENT OF PEDIATRIC PATIENTS (12 TO LESS THAN 18 YEARS OF AGE AND WEIGHING AT LEAST 30 KG) WITH PULMONARY MULTI-DRUG RESISTANT TUBERCULOSIS
- ODE-252 TREATMENT OF DUCHENE MUSCULAR DYSTROPHY IN PATIENTS 2 YEARS OF AGE TO LESS THAN 5 YEARS OF AGE
- ODE-253 INDICATED AS PART OF A COMBINATION REGIMEN WITH BEDAQUILINE AND LINEZOLID FOR THE TREATMENT OF ADULTS WITH PULMONARY EXTENSIVELY DRUG RESISTANT (XDR) OR TREATMENT-INTOLERANT OR NONRESPONSIVE MULTIDRUG-RESISTANT (MDR) TUBERCULOSIS (TB)
- ODE-254 INDICATED TO IMPROVE WAKEFULNESS IN ADULT PATIENTS WITH EXCESSIVE DAYTIME SLEEPINESS ASSOCIATED WITH NARCOLEPSY

PATENT AND EXCLUSIVITY TERMS**ORPHAN DRUG EXCLUSIVITY**

- ODE-255 INDICATED FOR THE TREATMENT OF EXCESSIVE DAYTIME SLEEPINESS (EDS) IN ADULT PATIENTS WITH NARCOLEPSY
- ODE-256 FOR HIV-1 INFECTION IN PEDIATRIC PTS AT LEAST 25 KG W/ NO ANTIRETROVIRAL (ARV) TX HX OR TO REPLACE CURRENT ARV REGIMEN FOR VIROLOGICALLY-SUPPRESSED ON STABLE ARV W/ NO HX TX FAILURE AND NO KNOWN SUBSTITUTIONS ASSOCIATED W/ RESISTANCE TO BIC, FTC, OR TAF
- ODE-257 IN COMBO W/ DEXAMETHASONE FOR ADULTS W/ RELAPSED OR REFRACTORY MULTIPLE MYELOMA WHO RECEIVED AT LEAST 4 PRIOR THERAPIES AND REFRACTORY TO AT LEAST 2 PROTEASOME INHIBITORS, AT LEAST 2 IMMUNOMODULATORY AGENTS, AND AN ANTI-CD38 MONOCLONAL ANTIBODY
- ODE-258 FOR THE TREATMENT OF CHRONIC HEPATITIS C VIRUS (HCV) GENOTYPE 2 OR 3 INFECTION IN PEDIATRIC PATIENTS BETWEEN 3 YEARS OF AGE AND 12 YEARS OF AGE OR WEIGHING 35 KG WITHOUT CIRRHOSIS OR WITH COMPENSATED CIRRHOSIS FOR USE IN COMBINATION WITH RIBAVIRIN
- ODE-259 INDICATED FOR THE TREATMENT OF ADULT PATIENTS WITH INTERMEDIATE-2 OR HIGH-RISK PRIMARY OR SECONDARY (POST-POLYCYTHEMIA VERA OR POST-ESSENTIAL THROMBOCYTHEMIA) MYELOFIBROSIS (MF)
- ODE-260 INDICATED TO INCREASE SYSTEMIC EXPOSURE OF ATAZANAVIR IN COMBINATION WITH OTHER ANTIRETROVIRAL AGENTS IN THE TREATMENT OF HIV-1 INFECTION IN PEDIATRIC PATIENTS WEIGHING AT LEAST 35 KG
- ODE-261 INDICATED TO SLOW THE RATE OF DECLINE IN PULMONARY FUNCTION IN PATIENTS WITH SYSTEMIC SCLEROSIS-ASSOCIATED INTERSTITIAL LUNG DISEASE (SSC-ILD)
- ODE-262 TREATMENT OF PEDIATRIC PATIENTS BETWEEN 3 YEARS OF AGE AND 12 YEARS OF AGE OR WEIGHING 35 KG WITH CHRONIC HEPATITIS C VIRUS (HCV) GENOTYPE 1, 4, 5, OR 6 INFECTION WITHOUT CIRRHOSIS OR WITH COMPENSATED CIRRHOSIS
- ODE-263 TREATMENT OF PEDIATRIC PATIENTS 3 YEARS OF AGE AND OLDER WITH CHRONIC HCV GENOTYPE 1 INFECTION WITH DECOMPENSATED CIRRHOSIS, FOR USE IN COMBINATION WITH RIBAVIRIN
- ODE-264 TREATMENT OF PEDIATRIC PATIENTS 3 YEARS OF AGE AND OLDER WITH CHRONIC HCV GENOTYPE 1 OR 4 INFECTION WHO ARE LIVER TRANSPLANT RECIPIENTS WITHOUT CIRRHOSIS OR WITH COMPENSATED CIRRHOSIS, FOR USE IN COMBINATION WITH RIBAVIRIN
- ODE-265 INDICATED FOR THE TREATMENT OF ADULT PATIENTS WITH METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) WHOSE TUMORS ARE ROS1-POSITIVE
- ODE-266 ADULT & PED >=12YRS OLD W/ SOLID TUMORS THAT HAVE NTRK W/O KNOWN ACQUIRED RESISTANCE MUTATION, ARE METASTATIC OR WHERE SURGICAL RESECTION IS LIKELY TO RESULT IN SEVERE MORBIDITY & HAVE EITHER PROGRESSED FOLLOWING TX OR HAVE NO SATISFACTORY ALTERNATIVE TX
- ODE-267 INDICATED IN COMBINATION WITH HIGH FLUID INTAKE, ALKALI, AND DIET MODIFICATION, FOR THE PREVENTION OF CYSTINE STONE FORMATION IN PEDIATRIC PATIENTS 20KG TO 9 YEARS OF AGE W/SEVERE HOMOZYGOUS CYSTINURIA, WHO ARE NOT RESPONSIVE TO THESE MEASURES ALONE
- ODE-268 INDICATED FOR TREATMENT OF PATIENTS WITH CUSHING'S DISEASE FOR WHOM PITUITARY SURGERY IS NOT AN OPTION OR HAS NOT BEEN CURATIVE
- ODE-269 PROPHYLAXIS OF ORGAN REJECTION IN PEDIATRIC PATIENTS RECEIVING ALLOGENEIC KIDNEY TRANSPLANT, LIVER TRANSPLANTS, AND HEART TRANSPLANT, IN COMBINATION WITH OTHER IMMUNOSUPPRESSANTS
- ODE-270 INDICATED TO INCREASE PAIN FREE LIGHT EXPOSURE IN ADULT PATIENTS WITH A HISTORY OF PHOTOTOXIC REACTIONS FROM ERYTHROPOIETIC PROTOPORPHYRIA (EPP)
- ODE-271 INDICATED IN COMBINATION WITH OTHER ANTI-MYELOMA PRODUCTS FOR THE TREATMENT OF ADULTS WITH MULTIPLE MYELOMA (MM)
- ODE-272 INDICATED FOR THE TREATMENT OF PULMONARY ARTERIAL HYPERTENSION (PAH) (WHO GROUP 1) TO DELAY DISEASE PROGRESSION
- ODE-273 INDICATED FOR THE TREATMENT OF ADULTS WITH ACUTE HEPATIC PORPHYRIA (AHP)
- ODE-274 INDICATED FOR THE TREATMENT OF ADULT PATIENTS WITH CHRONIC LYMPHOCYTIC LEUKEMIA (CLL) OR SMALL LYMPHOCYTIC LYMPHOMA (SLL)
- ODE-275 INDICATED FOR THE TREATMENT OF CYSTIC FIBROSIS (CF) IN PATIENTS AGED 12 YEARS AND OLDER WHO HAVE AT LEAST ONE F508DEL MUTATION IN THE CYSTIC FIBROSIS TRANSMEMBRANE CONDUCTANCE REGULATOR (CFTR) GENE
- ODE-276 INDICATED FOR THE TREATMENT OF ADULT PATIENTS WITH MANTLE CELL LYMPHOMA (MCL) WHO HAVE RECEIVED AT LEAST ONE PRIOR THERAPY

PATENT AND EXCLUSIVITY TERMS**ORPHAN DRUG EXCLUSIVITY**

- ODE-277 TX OF ADULTS W/ ADV OVARIAN, FALLOPIAN TUBE, OR PRIMARY PERITONEAL CANCER TREATED W/ >=3 PRIOR CHEMO REGIMENS & CANCER ASSOCIATED W/ HRD+ STATUS DEFINED BY A DELETERIOUS OR SUSPECTED DELETERIOUS BRCA MUTATION
- ODE-278 TX OF ADULTS W/ ADV OVARIAN, FALLOPIAN TUBE, OR PRIMARY PERITONEAL CANCER TREATED W/ >=3 PRIOR CHEMO REGIMENS & CANCER ASSOCIATED W/ HRD+ STATUS DEFINED BY GENOMIC INSTABILITY & PROGRESSED >6 MONTHS AFTER RESPONSE TO THE LAST PLATINUM-BASED CHEMOTHERAPY
- ODE-279 INDICATED FOR THE ACUTE TX OF INTERMITTENT, STEREOTYPIC EPISODES OF FREQUENT SEIZURE ACTIVITY (I.E. SEIZURE CLUSTERS, ACUTE REPETITIVE SEIZURES) THAT ARE DISTINCT FROM A PATIENT'S USUAL SEIZURE PATTERN IN PATIENTS WITH EPILEPSY 6 YEARS OF AGE AND OLDER
- ODE-280 INDICATED FOR THE TREATMENT OF DUCHENNE MUSCULAR DYSTROPHY (DMD) IN PATIENTS WHO HAVE A CONFIRMED MUTATION OF THE DMD GENE THAT IS AMENABLE TO EXON 53 SKIPPING
- ODE-281 INDICATED FOR THE TREATMENT OF SICKLE CELL DISEASE (SCD) IN ADULTS AND PEDIATRIC PATIENTS 12 YEARS OF AGE AND OLDER
- ODE-282 INDICATED TO SELECTIVELY STAIN THE INTERNAL LIMITING MEMBRANE (ILM)
- ODE-283 MAINTENANCE TX OF ADULTS W/ DELETERIOUS OR SUSPECTED DELETERIOUS GBRCA METASTATIC PANCREATIC ADENOCARCINOMA WHOSE DZ HAS NOT PROGRESSED ON >=16WKS OF 1ST LINE PLATINUM BASED CHEMO REGIMEN. SELECT PTS FOR THERAPY BASED ON APPROVED COMPANION DIAGNOSTIC
- ODE-284 IN COMBINATION WITH OTHER ANTIRETROVIRAL AGENTS, FOR THE TREATMENT OF HIV-1 INFECTION IN PEDIATRIC PATIENTS UNDER 12 YEARS OF AGE WEIGHING AT LEAST 35KG
- ODE-285 IN COMBINATION WITH OTHER ANTIRETROVIRAL AGENTS OTHER THAN PROTEASE INHIBITORS THAT REQUIRE A CYP3A INHIBITOR, FOR THE TREATMENT OF HIV-1 INFECTION IN PEDIATRIC PATIENTS UNDER 12 YEARS OF AGE WEIGHING AT LEAST 25KG AND LESS THAN 35KG
- ODE-286 INDICATED FOR THE TREATMENT OF ADULT PATIENTS WITH CUSHING'S DISEASE FOR WHOM PITUITARY SURGERY IS NOT AN OPTION OR HAS NOT BEEN CURATIVE
- ODE-287 TREATMENT OF ADULTS WITH NEWLY DIAGNOSED THERAPY-RELATED ACUTE MYELOID LEUKEMIA (T-AML) OR AML WITH MYELODYSPLASIA-RELATED CHANGES (AML-MRC)
- ODE-288 INDICATED FOR THE TREATMENT OF PEDIATRIC PATIENTS 2 YEARS OF AGE AND OLDER WITH NEUROFIBROMATOSIS TYPE 1 (NF1) WHO HAVE SYMPTOMATIC, INOPERABLE PLEXIFORM NEUROFIBROMAS (PN)
- ODE-289 INDICATED FOR THE TREATMENT OF ADULT PATIENTS WITH LOW-GRADE UPPER TRACT UROTHELIAL CANCER (LG-UTUC)
- ODE-290 INDICATED FOR THE INITIAL TREATMENT OF SEVERE MALARIA IN ADULT AND PEDIATRIC PATIENTS TO ALWAYS BE FOLLOWED BY A COMPLETE TREATMENT COURSE OF AN APPROPRIATE ORAL ANTIMALARIAL REGIMEN
- ODE-291 INDICATED FOR THE TREATMENT OF ADULT PATIENTS WITH METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) WHOSE TUMORS HAVE A MUTATION THAT LEADS TO MESENCHYMAL-EPITHELIAL TRANSITION (MET) EXON 14 SKIPPING AS DETECTED BY AN FDA-APPROVED TEST
- ODE-292 INDICATED FOR THE TREATMENT OF ADULTS WITH PREVIOUSLY TREATED, UNRESECTABLE LOCALLY ADVANCED OR METASTATIC CHOLANGIOCARCINOMA WITH A FIBROBLAST GROWTH FACTOR RECEPTOR 2 (FGFR2) FUSION OR OTHER REARRANGEMENT AS DETECTED BY AN FDA-APPROVED TEST
- ODE-293 TX OF PED PTS 6 YRS OF AGE & OLDER OR WEIGHING AT LEAST 17 KG WITH CHRONIC HEPATITIS C VIRUS GENOTYPE 1, 2, 3, 4, 5, OR 6 INFECTION: WITHOUT CIRRHOSIS OR WITH COMPENSATED CIRRHOSIS; OR WITH DECOMPENSATED CIRRHOSIS FOR USE IN COMBINATION WITH RIBAVIRIN
- ODE-294 PROPHYLAXIS OF ORGAN REJECTION IN PEDIATRIC PATIENTS RECEIVING ALLOGENEIC KIDNEY OR HEART TRANSPLANTS, IN COMBINATION WITH OTHER IMMUNOSUPPRESSANTS
- ODE-295 INDICATED FOR THE MAINTENANCE TREATMENT OF ADULT PATIENTS WITH ADVANCED EPITHELIAL OVARIAN CANCER WHO ARE IN A COMPLETE OR PARTIAL RESPONSE TO FIRST-LINE PLATINUM-BASED CHEMOTHERAPY
- ODE-296 INDICATED FOR THE TREATMENT OF ADULT PATIENTS WITH AIDS-RELATED KAPOSI SARCOMA (KS) AFTER FAILURE OF HIGHLY ACTIVE ANTIRETROVIRAL THERAPY (HAART)
- ODE-297 FOR THE TREATMENT OF KAPOSI SARCOMA (KS) IN ADULT PATIENTS WHO ARE HIV-NEGATIVE
- ODE-298 INDICATED FOR THE TREATMENT OF ADULT PATIENTS WITH ADVANCED GASTROINTESTINAL STROMAL TUMOR (GIST) WHO HAVE RECEIVED PRIOR TREATMENT WITH 3 OR MORE KINASE INHIBITORS, INCLUDING IMATINIB

PATENT AND EXCLUSIVITY TERMS**ORPHAN DRUG EXCLUSIVITY**

- ODE-299 INDICATED FOR THE TREATMENT OF ADULTS AND PEDIATRIC PATIENTS AGED 16 YEARS AND OLDER WITH METASTATIC OR LOCALLY ADVANCED EPITHELIOID SARCOMA NOT ELIGIBLE FOR COMPLETE RESECTION
- ODE-300 FOR THE TREATMENT OF ADULT PATIENTS WITH ANAPLASTIC LYMPHOMA KINASE (ALK)- POSITIVE METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) AS DETECTED BY AN FDA- APPROVED TEST, NOT INCLUDING PATIENTS WHO HAVE PROGRESSED ON OR ARE INTOLERANT TO CRIZOTINIB
- ODE-301 INDICATED FOR THE TREATMENT OF ADULT PATIENTS WITH METASTATIC RET FUSION- POSITIVE NON-SMALL CELL LUNG CANCER (NSCLC)
- ODE-302 INDICATED FOR THE TREATMENT OF ADULT AND PEDIATRIC PATIENTS 12 YEARS OF AGE AND OLDER WITH ADVANCED OR METASTATIC RET-MUTANT MEDULLARY THYROID CANCER (MTC) WHO REQUIRE SYSTEMIC THERAPY
- ODE-303 ADULT AND PEDIATRIC PATIENTS 12 YEARS OF AGE AND OLDER WITH ADVANCED OR METASTATIC RET FUSION-POSITIVE THYROID CANCER WHO REQUIRE SYSTEMIC THERAPY AND WHO ARE RADIOACTIVE IODINE-REFRACTORY (IF RADIOACTIVE IODINE IS APPROPRIATE)
- ODE-304 INDICATED FOR THE TREATMENT OF ADULT PATIENTS WITH METASTATIC SMALL CELL LUNG CANCER (SCLC) WITH DISEASE PROGRESSION ON OR AFTER PLATINUM-BASED CHEMOTHERAPY
- ODE-305 TREATMENT OF INVASIVE ASPERGILLOSIS
- ODE-306 W/ BEVACIZUMAB FOR MAINT TX OF ADULTS W/ ADV EPITHELIAL OVARIAN, FALLOPIAN TUBE, OR PRIMARY PERITONEAL CA IN COMPLETE OR PARTIAL RESPONSE TO 1ST LINE PT BASED CHEMO & WHOSE CA IS ASSOC W/ HOMOLOGOUS RECOMB DEF + STATUS DEFINED BY GENOMIC INSTABILITY
- ODE-307 INDICATED AS PART OF COMBINATION THERAPY IN THE TREATMENT OF PEDIATRIC PATIENTS 5 YEARS AND OLDER TO LESS THAN 12 YEARS OF AGE AND WEIGHING AT LEAST 15 KG WITH PULMONARY MULTI-DRUG RESISTANT TUBERCULOSIS (MDR-TB)
- ODE-308 TREATMENT OF PULMONARY ARTERIAL HYPERTENSION (PAH) (WHO GROUP 1) TO IMPROVE EXERCISE CAPACITY
- ODE-309 INDICATED FOR USE IN COMBINATION WITH TRASTUZUMAB AND CAPECITABINE FOR TREATMENT OF ADULT PATIENTS WITH METASTATIC HER2-POSITIVE BREAST CANCER AND BRAIN METASTASES, WHO HAVE RECEIVED ONE OR MORE PRIOR ANTI-HER2-BASED REGIMENS IN THE METASTATIC SETTING
- ODE-310 INDICATED FOR THE TREATMENT OF ADULT PATIENTS WITH RELAPSED OR REFRACTORY DIFFUSE LARGE B-CELL LYMPHOMA (DLBCL), NOT OTHERWISE SPECIFIED, INCLUDING DLBCL ARISING FROM FOLLICULAR LYMPHOMA, AFTER AT LEAST 2 LINES OF SYSTEMIC THERAPY
- ODE-311 INDICATED AS A SOURCE OF CALORIES AND FATTY ACIDS FOR THE TREATMENT OF PEDIATRIC AND ADULT PATIENTS WITH MOLECULARLY CONFIRMED LONG-CHAIN FATTY ACID OXIDATION DISORDERS (LC-FAOD)
- ODE-312 INDICATED FOR THE TREATMENT OF SEIZURES ASSOCIATED WITH DRAVET SYNDROME IN PATIENTS 2 YEARS OF AGE AND OLDER
- ODE-313 INDICATED FOR THE TREATMENT OF ADULT AND PEDIATRIC PATIENTS 12 YEARS OF AGE AND OLDER WITH SOLID TUMORS THAT HAVE A NEUROTROPHIC TYROSINE RECEPTOR KINASE (NTRK) GENE FUSION WITHOUT A KNOWN ACQUIRED RESISTANCE MUTATION, ARE METASTATIC OR WHERE SURGICAL RESECTION IS LIKELY TO RESULT IN SEVERE MORBIDITY, AND HAVE EITHER PROGRESSED FOLLOWING TREATMENT OR HAVE NO SATISFACTORY ALTERNATIVE THERAPY
- ODE-314 INDICATED FOR THE TREATMENT OF ADULT PATIENTS WITH RELAPSED OR REFRACTORY FOLLICULAR LYMPHOMA WHOSE TUMORS ARE POSITIVE FOR AN EZH2 MUTATION AS DETECTED BY AN FDA-APPROVED TEST AND WHO HAVE RECEIVED AT LEAST 2 PRIOR SYSTEMIC THERAPIES, AND FOR THE TREATMENT OF ADULT PATIENTS WITH RELAPSED OR REFRACTORY FOLLICULAR LYMPHOMA WHO HAVE NO SATISFACTORY ALTERNATIVE TREATMENT OPTIONS
- ODE-315 FOR CLEANSING OF THE COLON AS A PREPARATION FOR COLONOSCOPY IN PEDIATRIC PATIENTS 12 YEARS OF AGE AND OLDER
- ODE-316 TREATMENT OF ADULT PATIENTS WITH MYELODYSPLASTIC SYNDROMES (MDS), INCLUDING PREVIOUSLY TREATED AND UNTREATED, DE NOVO AND SECONDARY MDS WITH THE FOLLOWING FRENCH-AMERICAN-BRITISH SUBTYPES (REFRACTORY ANEMIA, REFRACTORY ANEMIA WITH RINGED SIDEROBLASTS, REFRACTORY ANEMIA WITH EXCESS BLASTS, AND CHRONIC MYELOMONOCYTIC LEUKEMIA [CMML]) AND INTERMEDIATE-1, INTERMEDIATE-2, AND HIGH-RISK INTERNATIONAL PROGNOSTIC SCORING SYSTEM GROUPS.
- ODE-317 FOR USE WITH POSITRON EMISSION TOMOGRAPHY (PET) FOR LOCALIZATION OF SOMATOSTATIN RECEPTOR POSITIVE NEUROENDOCRINE TUMORS (NETS) IN ADULT PATIENTS

PATENT AND EXCLUSIVITY TERMS**ORPHAN DRUG EXCLUSIVITY**

- ODE-318 TREATMENT OF ADULT PATIENTS WITH METASTATIC RET FUSION-POSITIVE NON-SMALL CELL LUNG CANCER (NSCLC) AS DETECTED BY AN FDA APPROVED TEST
- ODE-319 INDICATED IN PEDIATRIC PATIENTS (BIRTH TO LESS THAN 18 YEARS OF AGE AND WEIGHING AT LEAST 2.5 KG) FOR THE TREATMENT OF CHAGAS DISEASE (AMERICAN TRYPANOSOMIASIS) CAUSED BY TRYPANOSOMA CRUZI
- ODE-320 INDICATED FOR CONTINUED TREATMENT OF ADULT PATIENTS WITH ACUTE MYELOID LEUKEMIA WHO ACHIEVED FIRST COMPLETE REMISSION (CR) OR COMPLETE REMISSION WITH INCOMPLETE BLOOD COUNT RECOVERY (CRI) FOLLOWING INTENSIVE INDUCTION CHEMOTHERAPY AND ARE NOT ABLE TO COMPLETE INTENSIVE CURATIVE THERAPY
- ODE-321 TREATMENT OF CYSTIC FIBROSIS (CF) IN PATIENTS AGE 4 MONTHS TO LESS THAN 6 MONTHS WHO HAVE ONE MUTATION IN THE CFTR GENE THAT IS RESPONSIVE TO IVACAFTOR POTENTIATION BASED ON CLINICAL AND/OR IN VITRO ASSAY DATA
- ODE-322 TREATMENT OF NARCOLEPSY
- ODE-323 TREATMENT OF CYSTIC FIBROSIS (CF) IN PATIENTS AGED 12 YEARS AND OLDER WHO HAVE A MUTATION IN THE CYSTIC FIBROSIS TRANSMEMBRANE CONDUCTANCE REGULATOR (CFTR) GENE THAT IS RESPONSIVE BASED ON IN VITRO DATA
- ODE-324 TREATMENT OF HUTCHINSON-GILFORD PROGERIA SYNDROME (HGPS) AND PROGEROID LAMINOPATHIES
- ODE-325 TREATMENT OF ADULT PATIENTS WITH METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) HARBORING MESENCHYMAL-EPITHELIAL TRANSITION (MET) EXON 14 SKIPPING ALTERATIONS
- ODE-326 TREATMENT OF SEIZURES ASSOCIATED WITH LENNOX-GASTAUT SYNDROME (LGS) OR DRAVET SYNDROME (DS) IN PATIENTS BETWEEN 1 AND 2 YEARS OF AGE
- ODE-327 ADD-ON MAINTENANCE THERAPY TO IMPROVE PULMONARY FUNCTION IN ADULT PATIENTS 18 YEARS OF AGE AND OLDER WITH CYSTIC FIBROSIS AND WHO HAVE PASSED THE BRONCHITOL TOLERANCE TEST
- ODE-328 TREATMENT OF PEDIATRIC PATIENTS 1 YEAR OF AGE AND OLDER AND YOUNG ADULTS WITH RELAPSED OR REFRACTORY, SYSTEMIC ANAPLASTIC LARGE CELL LYMPHOMA (ALCL) THAT IS ALK-POSITIVE
- ODE-329 FOR THE TREATMENT OF NIGHTTIME SLEEP DISTURBANCES IN SMITH-MAGENIS SYNDROME (SMS) IN PEDIATRIC PATIENTS 3 TO 15 YEARS OF AGE
- ODE-330 THE TREATMENT OF NIGHTTIME SLEEP DISTURBANCES IN SMITH-MAGENIS SYNDROME (SMS) IN PATIENTS 16 YEARS OF AGE AND OLDER
- ODE-331 TREATMENT OF CATAPLEXY IN ADULT PATIENTS WITH NACROLEPSY
- ODE-332 TREATMENT OF SEIZURES ASSOCIATED WITH TUBEROUS SCLEROSIS COMPLEX (TSC) IN PATIENTS 1 YEAR OF AGE AND OLDER
- ODE-333 PROPHYLAXIS TO PREVENT ATTACKS OF HEREDITARY ANGIOEDEMA (HAE) IN ADULTS AND PEDIATRIC PATIENTS 12 YEARS OF AGE AND OLDER
- ODE-334 TREATMENT OF SPINAL MUSCULAR ATROPHY (SMA) IN PATIENTS 2 MONTHS OF AGE AND OLDER
- ODE-335 FOR TREATMENT OF CYSTIC FIBROSIS (CF) IN PATIENTS AGE 6 YEARS AND OLDER WHO HAVE AT LEAST ONE OF THE ADDITIONAL MUTATIONS IN THE CYSTIC FIBROSIS TRANSMEMBRANE CONDUCTANCE REGULATOR (CFTR) GENE THAT HAVE BEEN IDENTIFIED AS RESPONSIVE TO TEZACAFTOR/IVACAFTOR BASED ON IN VITRO DATA AND IDENTIFIED IN THE APPROVAL ON DECEMBER 21, 2020
- ODE-336 INDICATED FOR CHRONIC WEIGHT MANAGEMENT IN ADULT AND PEDIATRIC PATIENTS 6 YEARS OF AGE AND OLDER WITH OBESITY DUE TO PROOPIOMELANOCORTIN (POMC), PROPROTEIN CONVERTASE SUBTILISIN/KEXIN TYPE 1 (PCSK1), OR LEPTIN RECEPTOR (LEPR) DEFICIENCY CONFIRMED BY GENETIC TESTING DEMONSTRATING VARIANTS IN POMC, PCSK1, OR LEPR GENES THAT ARE INTERPRETED AS PATHOGENIC, LIKELY PATHOGENIC, OR OF UNCERTAIN SIGNIFICANCE (VUS)
- ODE-337 FOR ADJUVANT THERAPY AFTER TUMOR RESECTION IN ADULT PATIENTS WITH NON-SMALL CELL LUNG CANCER (NSCLC) WHOSE TUMORS HAVE EPIDERMAL GROWTH FACTOR RECEPTOR (EGFR) EXON 19 DELETIONS OR EXON 21 L858R MUTATIONS, AS DETECTED BY AN FDA-APPROVED TEST
- ODE-338 FOR THE TREATMENT OF CYSTIC FIBROSIS (CF) IN PATIENTS AGE 4 MONTHS AND OLDER WHO HAVE ONE OF THE ADDITIONAL MUTATIONS IN THE CYSTIC FIBROSIS TRANSMEMBRANE CONDUCTANCE REGULATOR (CFTR) GENE THAT HAVE BEEN IDENTIFIED AS RESPONSIVE TO IVACAFTOR POTENTIATION BASED ON IN VITRO DATA AND IDENTIFIED IN THE APPROVAL ON DECEMBER 21, 2020
- ODE-339 TREATMENT OF PRIMARY HYPEROXALURIA TYPE 1 (PH1) TO LOWER URINARY OXALATE LEVELS IN PEDIATRIC AND ADULT PATIENTS

PATENT AND EXCLUSIVITY TERMS**ORPHAN DRUG EXCLUSIVITY**

- ODE-340 TREATMENT OF ADULT AND PEDIATRIC PATIENTS 12 YEARS OF AGE AND OLDER WITH ADVANCED OR METASTATIC RET-MUTANT MEDULLARY THYROID CANCER (MTC) WHO REQUIRE SYSTEMIC THERAPY
- ODE-341 TREATMENT OF ADULT AND PEDIATRIC PATIENTS 12 YEARS OF AGE AND OLDER WITH ADVANCED OR METASTATIC RET FUSION-POSITIVE THYROID CANCER WHO REQUIRE SYSTEMIC THERAPY AND WHO ARE RADIOACTIVE IODINE-REFRACTORY (IF RADIOACTIVE IODINE IS APPROPRIATE)
- ODE-342 INDICATED TO REDUCE THE RISK OF MORTALITY IN PATIENTS WITH MOLYBDENUM COFACTOR DEFICIENCY (MOCD) TYPE A
- ODE-343 FOR TREATMENT OF ADULT PATIENTS WITH RELAPSED OR REFRACTORY MARGINAL ZONE LYMPHOMA (MZL) WHO HAVE RECEIVED AT LEAST ONE PRIOR ANTI-CD20-BASED REGIMEN
- ODE-344 FOR TREATMENT OF ADULT PATIENTS WITH RELAPSED OR REFRACTORY FOLLICULAR LYMPHOMA (FL) WHO HAVE RECEIVED AT LEAST THREE PRIOR LINES OF SYSTEMIC THERAPY
- ODE-345 IN PEDIATRIC AND ADULT PATIENTS AS ADJUNCTIVE THERAPY TO STANDARD OF CARE FOR THE TREATMENT OF ACUTE HYPERAMMONEMIA DUE TO PROPIONIC ACIDEMIA (PA) OR METHYLMALONIC ACIDEMIA (MMA)
- ODE-346 FOR THE TREATMENT OF ADULT PATIENTS WITH MULTIPLE MYELOMA WHO HAVE RECEIVED AT LEAST ONE PRIOR THERAPY, EXCLUDING ADULT PATIENTS COVERED BY XPOVIO'S PREVIOUS INDICATION FOR MULTIPLE MYELOMA APPROVED ON JULY 3, 2019
- ODE-347 TREATMENT OF DUCHENNE MUSCULAR DYSTROPHY (DMD) IN PATIENTS WHO HAVE A CONFIRMED MUTATION OF THE DMD GENE THAT IS AMENABLE TO EXON 45 SKIPPING
- ODE-348 TREATMENT OF ADULT PATIENTS WITH RELAPSED OR REFRACTORY MULTIPLE MYELOMA WHO HAVE RECEIVED AT LEAST FOUR PRIOR LINES OF THERAPY AND WHOSE DISEASE IS REFRACTORY TO AT LEAST ONE PROTEASOME INHIBITOR, ONE IMMUNOMODULATORY AGENT, AND ONE CD-38 DIRECTED MONOCLONAL ANTIBODY
- ODE-349 TREATMENT OF ADULT PATIENTS WITH METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) WHOSE TUMORS ARE ANAPLASTIC LYMPHOMA KINASE (ALK)-POSITIVE AS DETECTED BY AN FDA-APPROVED TEST, EXCLUDING PATIENTS WHOSE DISEASE HAS PROGRESSED ON CRIZOTINIB AND AT LEAST ONE OTHER ALK INHIBITOR FOR METASTATIC DISEASE; OR ALECTINIB OR CERITINIB AS THE FIRST ALK INHIBITOR THERAPY FOR METASTATIC DISEASE
- ODE-350 TREATMENT OF NEWLY-DIAGNOSED THERAPY-RELATED ACUTE MYELOID LEUKEMIA (T-AML) OR AML WITH MYELODYSPLASIA-RELATED CHANGES (AML-MRC) IN PEDIATRIC PATIENTS AGES 1 YEAR AND OLDER
- ODE-351 TREATMENT OF ADULT PATIENTS WITH PAROXYSMAL NOCTURNAL HEMOGLOBINURIA (PNH)
- ODE-352 TREATMENT OF ADULT PATIENTS WITH KRAS G12C-MUTATED LOCALLY ADVANCED OR METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC), AS DETERMINED BY AN FDA-APPROVED TEST, WHO HAVE RECEIVED AT LEAST ONE PRIOR SYSTEMIC THERAPY
- ODE-353 FOR TREATMENT OF ADULTS WITH PREVIOUSLY TREATED, UNRESECTABLE LOCALLY ADVANCED OR METASTATIC CHOLANGIOCARCINOMA WITH A FIBROBLAST GROWTH FACTOR RECEPTOR 2 (FGFR2) FUSION OR OTHER REARRANGEMENT AS DETECTED BY AN FDA-APPROVED TEST
- ODE-354 TREATMENT OF HUMAN SMALLPOX DISEASE CAUSED BY VARIOLA VIRUS IN ADULT AND PEDIATRIC PATIENTS, INCLUDING NEONATES
- ODE-355 FOR THE TREATMENT OF INVASIVE ASPERGILLOSIS IN ADULTS AND PEDIATRIC PATIENTS 13 YEARS OF AGE AND OLDER
- ODE-356 FOR THE TREATMENT OF ADULT PATIENTS WITH ADVANCED SYSTEMIC MASTOCYTOSIS (ADVSM). ADVSM INCLUDES PATIENTS WITH AGGRESSIVE SYSTEMIC MASTOCYTOSIS (ASM), SYSTEMIC MASTOCYTOSIS WITH AN ASSOCIATED HEMATOLOGICAL NEOPLASM (SM-AHN), AND MAST CELL LEUKEMIA (MCL)
- ODE-357 FOR THE TREATMENT OF CYSTIC FIBROSIS (CF) IN PATIENTS AGED 6 THROUGH 11 YEARS OLD WHO HAVE AT LEAST ONE F508DEL MUTATION IN THE CYSTIC FIBROSIS TRANSMEMBRANE CONDUCTANCE REGULATOR (CFTR) GENE OR A MUTATION IN THE CFTR GENE THAT IS RESPONSIVE BASED ON IN VITRO DATA
- ODE-358 FOR THE TREATMENT OF VERNAL KERATOCONJUNCTIVITIS (VKC) IN CHILDREN AND ADULTS
- ODE-359 FOR THE TREATMENT OF BOTH THE FIRST-STAGE (HEMOLYMPHATIC) AND SECOND-STAGE (MENINGOENCEPHALITIC) HUMAN AFRICAN TRYPAOSOMIASIS (HAT) DUE TO TRYPAOSOMA BRUCEI GAMBIENSE IN PATIENTS 6 YEARS OF AGE AND OLDER AND WEIGHING AT LEAST 20 KG
- ODE-360 FOR PROPHYLAXIS OF ORGAN REJECTION IN ADULT AND PEDIATRIC PATIENTS RECEIVING ALLOGENEIC LUNG TRANSPLANT
- ODE-361 INDICATED FOR THE TREATMENT OF CATAPLEXY OR EXCESSIVE DAYTIME SLEEPINESS (EDS)

PATENT AND EXCLUSIVITY TERMS**ORPHAN DRUG EXCLUSIVITY**

IN PATIENTS 7 YEARS OF AGE AND OLDER WITH NARCOLEPSY

- ODE-362 TREATMENT OF ADULT AND PEDIATRIC PATIENTS 12 YEARS AND OLDER WITH CHRONIC GRAFT-VERSUS-HOST DISEASE (CHRONIC GVHD) AFTER FAILURE OF AT LEAST TWO PRIOR LINES OF SYSTEMIC THERAPY
- ODE-363 TREATMENT OF PRURITUS IN PATIENTS 3 MONTHS OF AGE AND OLDER WITH PROGRESSIVE FAMILIAL INTRAHEPATIC CHOLESTASIS (PFIC)
- ODE-364 TREATMENT OF ADULT PATIENTS WITH VON HIPPEL-LINDAU (VHL) DISEASE WHO REQUIRE THERAPY FOR ASSOCIATED RENAL CELL CARCINOMA (RCC), CENTRAL NERVOUS SYSTEM (CNS) HEMANGIOBLASTOMAS, OR PANCREATIC NEUROENDOCRINE TUMORS (PNET), NOT REQUIRING IMMEDIATE SURGERY
- ODE-366 INDICATED FOR THE TREATMENT OF ADULTS WITH UNRESECTABLE OR METASTATIC GIST HARBORING A PLATELET-DERIVED GROWTH FACTOR RECEPTOR ALPHA (PDGFRA) EXON 18 MUTATION, INCLUDING PDGFRA D842V MUTATIONS
- ODE-367 PEDIATRIC PATIENTS AGED 6 MONTHS AND OLDER FOR THE TREATMENT OF C. DIFFICILE-ASSOCIATED DIARRHEA (CDAD)
- ODE-368 TREATMENT OF ADULT PATIENTS WITH PREVIOUSLY TREATED, LOCALLY ADVANCED OR METASTATIC CHOLANGIOCARCINOMA WITH AN ISOCITRATE DEHYDROGENASE-1 (IDH1) MUTATION AS DETECTED BY AN FDA-APPROVED TEST
- ODE-369 THE TREATMENT OF IDIOPATHIC HYPERSOMNIA (IH) IN ADULTS
- ODE-370 TREATMENT OF ADULT PATIENTS WITH RELAPSED OR REFRACTORY MARGINAL ZONE LYMPHOMA (MZL) WHO HAVE RECEIVED AT LEAST ONE ANTI-CD20-BASED REGIMEN
- ODE-371 TREATMENT OF ADULT PATIENTS WITH WALDENSTRM'S MACROGLOBULINEMIA (WM)
- ODE-372 FOR TREATMENT OF PEDIATRIC PATIENTS 3 YEARS OF AGE TO LESS THAN 12 YEARS OF AGE WEIGHING LESS THAN 45 KG WITH CHRONIC HEPATITIS C VIRUS (HCV) GENOTYPE 1, 2, 3, 4, 5 OR 6 INFECTION WITHOUT CIRRHOSIS OR WITH COMPENSATED CIRRHOSIS (CHILD-PUGH A); AND TREATMENT OF PEDIATRIC PATIENTS 3 YEARS OF AGE TO LESS THAN 12 YEARS OF AGE WEIGHING LESS THAN 45 KG WITH HCV GENOTYPE 1 INFECTION, WHO PREVIOUSLY HAVE BEEN TREATED WITH A REGIMEN CONTAINING AN HCV NS5A INHIBITOR OR AN NS3/4A PROTEASE INHIBITOR (PI), BUT NOT BOTH
- ODE-373 TREATMENT OF CHRONIC GRAFT-VERSUS-HOST DISEASE (CGVHD) AFTER FAILURE OF ONE OR TWO LINES OF SYSTEMIC THERAPY IN ADULT AND PEDIATRIC PATIENTS 12 YEARS AND OLDER
- ODE-374 TREATMENT OF ADULT PATIENTS WITH LOCALLY ADVANCED OR METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) WITH EPIDERMAL GROWTH FACTOR RECEPTOR (EGFR) EXON 20 INSERTION MUTATIONS, AS DETECTED BY AN FDA-APPROVED TEST, WHOSE DISEASE HAS PROGRESSED ON OR AFTER PLATINUM-BASED CHEMOTHERAPY
- ODE-375 THE TREATMENT OF ADULT AND PEDIATRIC PATIENTS 12 YEARS OF AGE AND OLDER WITH LOCALLY ADVANCED OR METASTATIC DIFFERENTIATED THYROID CANCER (DTC) THAT HAS PROGRESSED FOLLOWING PRIOR VEGFR-TARGETED THERAPY AND WHO ARE RADIOACTIVE IODINE-REFRACTORY OR INELIGIBLE
- ODE-376 FOR TREATMENT OF PEDIATRIC PATIENTS 3 YEARS OF AGE TO LESS THAN 6 YEARS OF AGE WEIGHING LESS THAN 17 KG WITH CHRONIC HEPATITIS C VIRUS (HCV) GENOTYPE 1, 2, 3, 4, 5, OR 6 INFECTION: WITHOUT CIRRHOSIS OR WITH COMPENSATED CIRRHOSIS; OR WITH DECOMPENSATED CIRRHOSIS FOR USE IN COMBINATION WITH RIBAVIRIN
- ODE-377 AS AN ADJUNCTIVE TREATMENT OF ADULT PATIENTS WITH SEVERE ACTIVE ANTI-NEUTROPHIL CYTOPLASMIC AUTOANTIBODY (ANCA)-ASSOCIATED VASCULITIS (GRANULOMATOSIS WITH POLYANGIITIS [GPA] AND MICROSCOPIC POLYANGIITIS [MPA])
- ODE-378 A COMPLETE REGIMEN FOR THE TREATMENT OF HUMAN IMMUNODEFICIENCY VIRUS TYPE 1 (HIV-1) INFECTION IN PEDIATRIC PATIENTS WEIGHING 14 KG TO LESS THAN 25 KG WHO HAVE NO ANTIRETROVIRAL TREATMENT HISTORY OR TO REPLACE THE CURRENT ANTIRETROVIRAL REGIMEN IN THOSE WHO ARE VIROLOGICALLY-SUPPRESSED (HIV-1 RNA LESS THAN 50 COPIES PER ML) ON A STABLE ANTIRETROVIRAL REGIMEN WITH NO HISTORY OF TREATMENT FAILURE AND NO KNOWN SUBSTITUTIONS ASSOCIATED WITH RESISTANCE TO THE INDIVIDUAL COMPONENTS OF BIKTARVY
- ODE-379 FOR TREATMENT OF CHOLESTATIC PRURITUS IN PATIENTS WITH ALAGILLE SYNDROME (ALGS) 1 YEAR OF AGE AND OLDER
- ODE-380 TREATMENT OF PEDIATRIC PATIENTS GREATER THAN OR EQUAL TO 1 YEAR OF AGE WITH ACCELERATED PHASE PHILADELPHIA CHROMOSOME POSITIVE CHRONIC MYELOID LEUKEMIA (PH+ CML) WITH RESISTANCE OR INTOLERANCE TO PRIOR TYROSINE-KINASE INHIBITOR (TKI) THERAPY
- ODE-381 TREATMENT OF ADULT PATIENTS WITH PHILADELPHIA CHROMOSOME-POSITIVE CHRONIC MYELOID LEUKEMIA (PH+ CML) IN CHRONIC PHASE (CP) WITH THE T315I MUTATION

PATENT AND EXCLUSIVITY TERMS**ORPHAN DRUG EXCLUSIVITY**

- ODE-382 TREATMENT OF ADULT PATIENTS WITH PHILADELPHIA CHROMOSOME-POSITIVE CHRONIC MYELOID LEUKEMIA (PH+ CML) IN CHRONIC PHASE (CP), PREVIOUSLY TREATED WITH TWO OR MORE TYROSINE KINASE INHIBITORS (TKIS)
- ODE-383 FOR USE WITH POSITRON EMISSION TOMOGRAPHY (PET) FOR THE LOCALIZATION OF KNOWN SOMATOSTATIN RECEPTOR POSITIVE NEUROENDOCRINE TUMORS (NETS) IN ADULT AND PEDIATRIC PATIENTS
- ODE-384 TO REDUCE THE RISK OF OTOTOXICITY ASSOCIATED WITH CISPLATIN IN PEDIATRIC PATIENTS 1 MONTH OF AGE AND OLDER WITH LOCALIZED, NON-METASTATIC SOLID TUMORS
- ODE-385 FOR TREATMENT OF ENDOGENOUS HYPERCORTISOLEMIA IN ADULT PATIENTS WITH CUSHING'S SYNDROME FOR WHOM SURGERY IS NOT AN OPTION OR HAS NOT BEEN CURATIVE
- ODE-386 FOR THE TREATMENT OF ADULT PATIENTS WITH LOCALLY ADVANCED UNRESECTABLE OR METASTATIC MALIGNANT PERIVASCULAR EPITHELIOID CELL TUMOR (PECOMA)
- ODE-387 TO INCREASE LINEAR GROWTH IN PEDIATRIC PATIENTS WITH ACHONDROPLASIA WHO ARE 5 YEARS OF AGE AND OLDER WITH OPEN EPIPHYSES
- ODE-388 FOR THE TREATMENT OF ADULTS AND PEDIATRIC PATIENTS (12 YEARS OF AGE AND OLDER AND WEIGHING AT LEAST 35 KG) WITH POST-TRANSPLANT CYTOMEGALOVIRUS (CMV) INFECTION/DISEASE THAT IS REFRACTORY TO TREATMENT (WITH OR WITHOUT GENOTYPIC RESISTANCE) WITH GANCICLOVIR, VALGANCICLOVIR, CIDOFOVIR OR FOSCARNET
- ODE-389 TO REDUCE PROTEINURIA IN ADULTS WITH PRIMARY IMMUNOGLOBULIN A NEPHROPATHY (IGAN) AT RISK OF RAPID DISEASE PROGRESSION, GENERALLY A URINE PROTEIN-TO-CREATININE RATIO (UPCR) > OR = 1.5 G/G
- ODE-390 AS AN ADJUNCT FOR INTRAOPERATIVE IDENTIFICATION OF MALIGNANT LESIONS IN ADULT PATIENTS WITH OVARIAN CANCER
- ODE-391 TREATMENT OF FACIAL ANGIOFIBROMA ASSOCIATED WITH TUBEROUS SCLEROSIS IN ADULTS AND PEDIATRIC PATIENTS 6 YEARS OF AGE AND OLDER
- ODE-392 TREATMENT OF HEMOLYTIC ANEMIA IN ADULTS WITH PYRUVATE KINASE (PK) DEFICIENCY
- ODE-393 TREATMENT OF SEIZURES ASSOCIATED WITH LENNOX-GASTAUT SYNDROME (LGS) IN PATIENTS 2 YEARS OF AGE AND OLDER
- ODE-394 FOR TREATMENT OF SICKLE CELL DISEASE (SCD) IN PEDIATRIC PATIENTS 4 YEARS OF AGE TO LESS THAN 12 YEARS OF AGE
- ODE-395 TREATMENT OF SEIZURES ASSOCIATED WITH CYCLIN-DEPENDENT KINASE-LIKE 5 (CDKL5) DEFICIENCY DISORDER (CDD) IN PATIENTS 2 YEARS OF AGE AND OLDER
- ODE-396 TREATMENT OF ADULT AND PEDIATRIC PATIENTS 2 YEARS OF AGE AND OLDER WITH SEVERE MANIFESTATIONS OF PIK3CA-RELATED OVERGROWTH SPECTRUM (PROS) WHO REQUIRE SYSTEMIC THERAPY
- ODE-397 TREATMENT OF ADULTS WITH INTERMEDIATE OR HIGH-RISK PRIMARY OR SECONDARY (POST-POLYCYTHEMIA VERA OR POST-ESSENTIAL THROMBOCYTHEMIA) MYELOFIBROSIS (MF) WITH A PLATELET COUNT BELOW $50 \times 10^9/L$
- ODE-398 TREATMENT OF ADULTS WITH SYMPTOMATIC NEW YORK HEART ASSOCIATION (NYHA) CLASS II-III OBSTRUCTIVE HYPERTROPHIC CARDIOMYOPATHY (HCM) TO IMPROVE FUNCTIONAL CAPACITY AND SYMPTOMS
- ODE-399 TREATMENT OF PEDIATRIC PATIENTS AGED ONE MONTH AND OLDER WITH NEWLY DIAGNOSED JUVENILE MYELOMONOCYTIC LEUKEMIA (JMML)
- ODE-400 TREATMENT OF SPINAL MUSCULAR ATROPHY (SMA) IN PEDIATRIC PATIENTS BETWEEN BIRTH AND 2 MONTHS OF AGE
- ODE-401 TREATMENT OF ADULT PATIENTS WITH STABLE WILSON'S DISEASE WHO ARE DE-COPPERED AND TOLERANT TO PENICILLAMINE
- ODE-402 FOR CHRONIC WEIGHT MANAGEMENT IN ADULT AND PEDIATRIC PATIENTS 6 YEARS OF AGE AND OLDER WITH MONOGENIC OR SYNDROMIC OBESITY DUE TO BARDET-BIEDL SYNDROME (BBS)
- ODE-403 TREATMENT OF SEIZURES ASSOCIATED WITH DRAVET SYNDROME (DS) IN PATIENTS TAKING CLOBAZAM WHO ARE 6 MONTHS TO LESS THAN 2 YEARS OF AGE AND WEIGHING 7 KG OR MORE
- ODE-404 TREATMENT OF ADULTS WITH RELAPSED OR REFRACTORY MYELOID/LYMPHOID NEOPLASMS (MLNS) WITH FIBROBLAST GROWTH FACTOR RECEPTOR 1 (FGFR1) REARRANGEMENT
- ODE-405 TREATMENT OF PEDIATRIC PATIENTS AGE 1 YEAR AND OLDER WITH CHRONIC GRAFT-VERSUS-HOST DISEASE (CGVHD) AFTER FAILURE OF ONE OR MORE LINES OF SYSTEMIC THERAPY
- ODE-406 TO IMPROVE KIDNEY FUNCTION IN ADULTS WITH HEPATORENAL SYNDROME WITH RAPID REDUCTION IN KIDNEY FUNCTION
- ODE-407 TREATMENT OF ADULT AND PEDIATRIC PATIENTS 1 YEAR OF AGE AND OLDER WITH UNRESECTABLE, RECURRENT, OR REFRACTORY INFLAMMATORY MYOFIBROBLASTIC TUMOR (IMT)

PATENT AND EXCLUSIVITY TERMS**ORPHAN DRUG EXCLUSIVITY**

THAT IS ALK-POSITIVE

- ODE-408 TREATMENT OF CYSTIC FIBROSIS (CF) IN PATIENTS AGED 1 YEAR TO LESS THAN 2 YEARS WHO ARE HOMOZYGOUS FOR THE F508DEL MUTATION IN THE CFTR GENE
- ODE-409 TREATMENT OF ADULT PATIENTS WITH LOCALLY ADVANCED OR METASTATIC SOLID TUMORS WITH A RET GENE FUSION THAT HAVE PROGRESSED ON OR FOLLOWING PRIOR SYSTEMIC TREATMENT OR WHO HAVE NO SATISFACTORY ALTERNATIVE TREATMENT OPTIONS
- ODE-410 TREATMENT OF ADULT PATIENTS WITH PREVIOUSLY TREATED, UNRESECTABLE, LOCALLY ADVANCED OR METASTATIC INTRAHEPATIC CHOLANGIOCARCINOMA HARBORING FIBROBLAST GROWTH FACTOR RECEPTOR 2 (FGFR2) GENE FUSIONS OR OTHER REARRANGEMENTS
- ODE-411 TREATMENT OF AMYOTROPHIC LATERAL SCLEROSIS (ALS) IN ADULTS
- ODE-412 TREATMENT OF ADULT PATIENTS WITH LOCALLY ADVANCED NON-SMALL CELL LUNG CANCER (NSCLC) WITH A REARRANGED DURING TRANSFECTION (RET) GENE FUSION, AS DETECTED BY AN FDA-APPROVED TEST
- ODE-413 TREATMENT OF ADULT PATIENTS WITH RELAPSED OR REFRACTORY ACUTE MYELOID LEUKEMIA (AML) WITH A SUSCEPTIBLE ISOCITRATE DEHYDROGENASE-1 (IDH1) MUTATION AS DETECTED BY AN FDA-APPROVED TEST
- ODE-414 TREATMENT OF NEONATAL SEIZURES IN TERM AND PRETERM INFANTS
- ODE-415 TREATMENT OF PRIMARY HYPEROXALURIA TYPE 1 (PH1) TO LOWER PLASMA OXALATE LEVELS IN PEDIATRIC AND ADULT PATIENTS
- ODE-416 TREATMENT OF ADULT PATIENTS WITH HISTIOCYTIC NEOPLASMS
- ODE-417 TREATMENT OF TRANSFUSIONAL IRON OVERLOAD IN ADULT PATIENTS WITH THALASSEMIA SYNDROMES EXCLUDING ADULT PATIENTS COVERED BY THE INDICATION FOR THALASSEMIA SYNDROMES APPROVED ON OCTOBER 14, 2011
- ODE-418 TREATMENT OF TRANSFUSIONAL IRON OVERLOAD IN PEDIATRIC PATIENTS 3 YEARS OF AGE AND OLDER WITH THALASSEMIA SYNDROMES
- ODE-419 TREATMENT OF TRANSFUSIONAL IRON OVERLOAD IN ADULT AND PEDIATRIC PATIENTS 3 YEARS OF AGE AND OLDER WITH SICKLE CELL DISEASE OR OTHER ANEMIAS
- ODE-420 TREATMENT OF TRANSFUSIONAL IRON OVERLOAD IN PEDIATRIC PATIENTS 8 YEARS OF AGE AND OLDER WITH THALASSEMIA SYNDROMES
- ODE-421 TREATMENT OF TRANSFUSIONAL IRON OVERLOAD IN ADULT AND PEDIATRIC PATIENTS 8 YEARS OF AGE AND OLDER WITH SICKLE CELL DISEASE OR OTHER ANEMIAS
- ODE-422 TREATMENT OF ADULT PATIENTS WITH RAS WILD-TYPE, HER2-POSITIVE UNRESECTABLE OR METASTATIC COLORECTAL CANCER THAT HAS PROGRESSED FOLLOWING TREATMENT WITH FLUOROPYRIMIDINE-, OXALIPLATIN-, AND IRINOTECAN-BASED CHEMOTHERAPY
- ODE-423 FOR PROPHYLAXIS OF CYTOMEGALOVIRUS (CMV) DISEASE IN ADULT KIDNEY TRANSPLANT RECIPIENTS AT HIGH RISK (DONOR CMV SEROPOSITIVE/RECIPIENT CMV SERONEGATIVE [D+/R-])
- ODE-424 TREATMENT OF ADULT PATIENTS WITH RELAPSED OR REFRACTORY MANTLE CELL LYMPHOMA (MCL) AFTER AT LEAST TWO LINES OF SYSTEMIC THERAPY, INCLUDING A BTK INHIBITOR
- ODE-425 TREATMENT OF RETT SYNDROME IN ADULTS AND PEDIATRIC PATIENTS 2 YEARS OF AGE AND OLDER
- ODE-426 FOR USE IN PATIENTS 18 YEARS OF AGE OR OLDER WHO HAVE LIMITED OR NO ALTERNATIVE OPTIONS FOR THE TREATMENT OF CANDIDEMIA AND INVASIVE CANDIDIASIS
- ODE-427 TREATMENT OF FRIEDREICH'S ATAXIA IN ADULTS AND ADOLESCENTS AGED 16 YEARS AND OLDER
- ODE-428 TREATMENT OF PEDIATRIC PATIENTS 1 YEAR OF AGE AND OLDER WITH LOW-GRADE GLIOMA (LGG) WITH A BRAF V600E MUTATION WHO REQUIRE SYSTEMIC THERAPY
- ODE-429 TREATMENT OF CHOLESTATIC PRURITUS IN PATIENTS 3 MONTHS OF AGE TO LESS THAN 1 YEAR OF AGE WITH ALAGILLE SYNDROME (ALGS)
- ODE-430 TREATMENT OF ACTIVATED PHOSPHOINOSITIDE 3-KINASE DELTA (PI3K DELTA) SYNDROME (APDS) IN ADULT AND PEDIATRIC PATIENTS 12 YEARS OF AGE AND OLDER
- ODE-431 TREATMENT OF CATAPLEXY OR EXCESSIVE DAYTIME SLEEPINESS (EDS) IN ADULTS WITH NARCOLEPSY
- ODE-432 TREATMENT OF AMYOTROPHIC LATERAL SCLEROSIS (ALS) IN ADULTS WHO HAVE A MUTATION IN THE SUPEROXIDE DISMUTASE 1 (SOD1) GENE
- ODE-433 TREATMENT OF CYSTIC FIBROSIS (CF) IN PATIENTS AGED 2 YEARS TO LESS THAN 6 YEARS WHO HAVE AT LEAST ONE F508DEL MUTATION IN THE CYSTIC FIBROSIS TRANSMEMBRANE CONDUCTANCE REGULATOR (CFTR) GENE OR A MUTATION IN THE CFTR GENE THAT IS

PATENT AND EXCLUSIVITY TERMS

ADB 72 of 225

ORPHAN DRUG EXCLUSIVITY

RESPONSIVE BASED ON IN VITRO DATA

- ODE-434 TREATMENT OF ADULT PATIENTS WITH INDOLENT SYSTEMIC MASTOCYTOSIS (ISM)
- ODE-435 THE TREATMENT OF CYSTIC FIBROSIS (CF) IN PATIENTS 1 MONTH TO LESS THAN 4 MONTHS OF AGE WHO HAVE AT LEAST ONE MUTATION IN THE CFTR GENE THAT IS RESPONSIVE TO IVACAFTOR POTENTIATION BASED ON CLINICAL AND/OR IN VITRO ASSAY DATA
- ODE-436 TREATMENT OF CHOLESTATIC PRURITUS IN PATIENTS 12 MONTHS OF AGE AND OLDER WITH ALAGILLE SYNDROME (ALGS)
- ODE-437 FOR USE IN COMBINATION WITH STANDARD INDUCTION AND CONSOLIDATION, AND AS MAINTENANCE THERAPY FOLLOWING CONSOLIDATION CHEMOTHERAPY, FOR THE TREATMENT OF ADULT PATIENTS WITH NEWLY DIAGNOSED ACUTE MYELOID LEUKEMIA (AML) THAT IS FLT3 INTERNAL TANDEM DUPLICATION (ITD)-POSITIVE AS DETECTED BY AN FDA-APPROVED TEST
- ODE-438 AS A LIVER-DIRECTED TREATMENT FOR ADULT PATIENTS WITH UVEAL MELANOMA WITH UNRESECTABLE HEPATIC METASTASES AFFECTING LESS THAN 50% OF THE LIVER AND NO EXTRAHEPATIC DISEASE OR EXTRAHEPATIC DISEASE LIMITED TO THE BONE, LYMPH NODES, SUBCUTANEOUS TISSUES, OR LUNG THAT IS AMENABLE TO RESECTION OR RADIATION
- ODE-439 FOR THE REDUCTION IN VOLUME OF NEW HETEROTOPIC OSSIFICATION IN ADULTS AND PEDIATRIC PATIENTS AGED 8 YEARS AND OLDER FOR FEMALES AND 10 YEARS AND OLDER FOR MALES WITH FIBRODYSPLASIA OSSIFICANS PROGRESSIVA (FOP)
- ODE-440 FOR TREATMENT OF ADULTS WITH CHOREA ASSOCIATED WITH HUNTINGTON'S DISEASE
- ODE-441 TREATMENT OF INTERMEDIATE OR HIGH-RISK MYELOFIBROSIS (MF), INCLUDING PRIMARY MF OR SECONDARY MF [POST-POLYCYTHEMIA VERA (PV) AND POST-ESSENTIAL THROMBOCYTHEMIA (ET)], IN ADULTS WITH ANEMIA
- ODE-442 TO MOBILIZE HEMATOPOIETIC STEM CELLS TO THE PERIPHERAL BLOOD FOR COLLECTION AND SUBSEQUENT AUTOLOGOUS TRANSPLANTATION IN PATIENTS WITH MULTIPLE MYELOMA
- ODE-443 TO LOWER URINARY OXALATE LEVELS IN CHILDREN 9 YEARS OF AGE AND OLDER AND ADULTS WITH PRIMARY HYPEROXALURIA TYPE 1 (PH1) AND RELATIVELY PRESERVED KIDNEY FUNCTION, E.G., EGFR GREATER THAN OR EQUAL TO 30 ML/MIN/1.73 M²
- ODE-444 TREATMENT OF PEDIATRIC PATIENTS 1 YEAR OF AGE AND OLDER WITH CHRONIC PHASE (CP) PHILADELPHIA CHROMOSOME-POSITIVE CHRONIC MYELOGENOUS LEUKEMIA (PH+ CML), NEWLY-DIAGNOSED OR RESISTANT OR INTOLERANT TO PRIOR THERAPY
- ODE-445 TREATMENT OF ADULT PATIENTS WITH METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) WITH A BRAF V600E MUTATION, AS DETECTED BY AN FDA-APPROVED TEST
- ODE-446 TREATMENT OF GENERALIZED MYASTHENIA GRAVIS (GMG) IN ADULT PATIENTS WHO ARE ANTI-ACETYLCHOLINE RECEPTOR (ACHR) ANTIBODY POSITIVE
- ODE-447 TREATMENT OF ADULT PATIENTS WITH RELAPSED OR REFRACTORY MYELODYSPLASTIC SYNDROMES (MDS) WITH A SUSCEPTIBLE ISOCITRATE DEHYDROGENASE-1 (IDH1) MUTATION AS DETECTED BY AN FDA-APPROVED TEST
- ODE-448 TREATMENT OF PEDIATRIC PATIENTS OLDER THAN 1 MONTH UP TO 12 YEARS OF AGE WITH SOLID TUMORS THAT HAVE A NEUROTROPHIC TYROSINE RECEPTOR KINASE (NTRK) GENE FUSION, AS DETECTED BY AN FDA-APPROVED TEST WITHOUT A KNOWN ACQUIRED RESISTANCE MUTATION, ARE METASTATIC OR WHERE SURGICAL RESECTION IS LIKELY TO RESULT IN SEVERE MORBIDITY, AND HAVE PROGRESSED FOLLOWING TREATMENT OR HAVE NO SATISFACTORY ALTERNATIVE THERAPY
- ODE-449 TO INCREASE LINEAR GROWTH IN PEDIATRIC PATIENTS LESS THAN 5 YEARS OF AGE WITH ACHONDROPLASIA WITH OPEN EPIPHYSES
- ODE-450 TREATMENT OF DUCHENNE MUSCULAR DYSTROPHY (DMD) IN PATIENTS 2 YEARS OF AGE AND OLDER
- ODE-451 TREATMENT OF ADULT PATIENTS WITH CHRONIC LYMPHOCYTIC LEUKEMIA OR SMALL LYMPHOCYTIC LYMPHOMA (CLL/SLL) WHO HAVE RECEIVED AT LEAST TWO PRIOR LINES OF THERAPY, INCLUDING A BTK INHIBITOR AND A BCL-2 INHIBITOR
- ODE-452 FOR ADULT PATIENTS WITH PROGRESSING DESMOID TUMORS WHO REQUIRE SYSTEMIC TREATMENT
- ODE-453 TREATMENT OF INVASIVE MUCORMYCOSIS IN PEDIATRIC PATIENTS 1 YEAR OF AGE AND OLDER
- ODE-454 TREATMENT OF INVASIVE MUCORMYCOSIS IN PEDIATRIC PATIENTS 6 YEARS OF AGE AND OLDER WHO WEIGH 16 KG AND GREATER
- ODE-455 TREATMENT OF ADULT PATIENTS WITH LOCALLY ADVANCED OR METASTATIC ROS1-POSITIVE NON-SMALL CELL LUNG CANCER (NSCLC) WITH ADENOCARCINOMA HISTOLOGY
- ODE-456 TREATMENT OF ADULTS WITH PAROXYSMAL NOCTURNAL HEMOGLOBINURIA (PNH)

PATENT AND EXCLUSIVITY TERMS**ORPHAN DRUG EXCLUSIVITY**

- ODE-457 TREATMENT OF HIV-1 INFECTION IN PEDIATRIC PATIENTS UNDER 12 YEARS OF AGE WEIGHING AT LEAST 14 KG TO LESS THAN 25 KG
- ODE-458 TREATMENT OF INVASIVE ASPERGILLOSIS IN PEDIATRIC PATIENTS 6 YEARS OF AGE AND OLDER WHO WEIGH 16 KILOGRAMS (KG) AND GREATER
- ODE-459 TREATMENT OF INVASIVE ASPERGILLOSIS IN PEDIATRIC PATIENTS 1 YEAR OF AGE AND OLDER
- ODE-460 TREATMENT OF WOUNDS ASSOCIATED WITH DYSTROPHIC AND JUNCTIONAL EPIDERMOLYSIS BULLOSA (EB) IN ADULT AND PEDIATRIC PATIENTS 6 MONTHS OF AGE AND OLDER
- ODE-461 TREATMENT OF THE POLYNEUROPATHY OF HEREDITARY TRANSTHYRETIN-MEDIATED AMYLOIDOSIS IN ADULTS
- ODE-462 TO REDUCE THE RISK OF RELAPSE IN ADULT AND PEDIATRIC PATIENTS WITH HIGH-RISK NEUROBLASTOMA (HRNB) WHO HAVE DEMONSTRATED AT LEAST A PARTIAL RESPONSE TO PRIOR MULTIAGENT, MULTIMODALITY THERAPY INCLUDING ANTI-GD2 IMMUNOTHERAPY
- ODE-463 FOR FIRST-LINE TREATMENT OF ADULT PATIENTS WITH METASTATIC PANCREATIC ADENOCARCINOMA
- ODE-464 TO REDUCE THE LOSS OF KIDNEY FUNCTION IN ADULTS WITH PRIMARY IMMUNOGLOBULIN A NEPHROPATHY (IGAN) WHO ARE AT RISK FOR DISEASE PROGRESSION, EXCLUDING THE USE PROVIDED FOR IN THE INDICATION APPROVED ON DECEMBER 15, 2021
- ODE-465 TREATMENT OF SEVERE FROSTBITE IN ADULTS TO REDUCE THE RISK OF DIGIT AMPUTATIONS
- ODE-466 TREATMENT OF ADULT AND PEDIATRIC PATIENTS 11 YEARS OF AGE AND OLDER WITH EOSINOPHILIC ESOPHAGITIS (EOE)
- ODE-467 TREATMENT OF ADULT PATIENTS WITH RELAPSED OR REFRACTORY FOLLICULAR LYMPHOMA (FL), AFTER TWO OR MORE LINES OF SYSTEMIC THERAPY
- ODE-468 TREATMENT OF HUMAN IMMUNODEFICIENCY VIRUS TYPE 1 (HIV-1) INFECTION IN PEDIATRIC PATIENTS WEIGHING AT LEAST 14 KG TO REPLACE THE CURRENT ANTIRETROVIRAL REGIMEN IN THOSE WHO ARE VIROLOGICALLY SUPPRESSED (HIV-1 RNA LESS THAN 50 COPIES PER ML) ON A STABLE ANTIRETROVIRAL REGIMEN WITH NO KNOWN OR SUSPECTED SUBSTITUTIONS ASSOCIATED WITH RESISTANCE TO BICTEGRAVIR OR TENOFOVIR AND WITH KNOWN OR SUSPECTED SUBSTITUTIONS ASSOCIATED WITH RESISTANCE TO EMTRICITABINE
- ODE-469 TREATMENT OF PULMONARY ARTERIAL HYPERTENSION (PAH) (WHO GROUP I) IN PEDIATRIC PATIENTS 1 TO 17 YEARS OLD TO IMPROVE EXERCISE ABILITY AND, IN PEDIATRIC PATIENTS TOO YOUNG TO PERFORM STANDARDIZED EXERCISE TESTING, PULMONARY HEMODYNAMICS THOUGHT TO UNDERLY IMPROVEMENTS IN EXERCISE
- ODE-470 TO REDUCE THE FREQUENCY OF PAINFUL CRISES AND REDUCE THE NEED FOR BLOOD TRANSFUSIONS IN PEDIATRIC PATIENTS AGED 6 MONTHS OF AGE TO LESS THAN 2 YEARS, WITH SICKLE CELL ANEMIA WITH RECURRENT MODERATE TO SEVERE PAINFUL CRISES
- ODE-471 TREATMENT OF CHOLESTATIC PRURITUS IN PATIENTS 5 YEARS OF AGE AND OLDER WITH PROGRESSIVE FAMILIAL INTRAHEPATIC CHOLESTASIS (PFIC)
- ODE-472 TREATMENT OF ADULT PATIENTS WITH NEWLY DIAGNOSED PHILADELPHIA CHROMOSOME-POSITIVE ACUTE LYMPHOBLASTIC LEUKEMIA (PH+ ALL)
- ODE-473 TREATMENT OF DUCHENNE MUSCULAR DYSTROPHY (DMD) IN PATIENTS 6 YEARS OF AGE AND OLDER
- ODE-474 LONG-TERM MAINTENANCE TREATMENT IN ACROMEGALY PATIENTS WHO HAVE RESPONDED TO AND TOLERATED TREATMENT WITH OCTREOTIDE OR LANREOTIDE
- ODE-475 CHRONIC TREATMENT OF ADULTS WITH PULMONARY ARTERIAL HYPERTENSION (PAH, WHO GROUP I AND WHO FUNCTIONAL CLASS (FC) II-III)
- ODE-476 TREATMENT OF EXTRAVASCULAR HEMOLYSIS (EVH) IN ADULTS WITH PAROXYSMAL NOCTURNAL HEMOGLOBINURIA (PNH)
- ODE-477 ADJUVANT TREATMENT IN ADULT PATIENTS FOLLOWING TUMOR RESECTION OF ANAPLASTIC LYMPHOMA KINASE (ALK)-POSITIVE NON-SMALL CELL LUNG CANCER (NSCLC) (TUMORS GREATER THAN OR EQUAL TO 4 CM OR NODE POSITIVE), AS DETECTED BY AN FDA-APPROVED TEST
- ODE-478 TREATMENT OF PATIENTS 6 MONTHS OF AGE AND OLDER WITH RELAPSED OR REFRACTORY PEDIATRIC LOW-GRADE GLIOMA (LGG) HARBORING A BRAF FUSION OR REARRANGEMENT, OR BRAF V600 MUTATION
- ODE-479 TREATMENT OF PEDIATRIC PATIENTS 12 YEARS AND OLDER WITH SOMATOSTATIN RECEPTOR-POSITIVE GASTROENTEROPANCREATIC NEUROENDOCRINE TUMORS (GEP-NETS), INCLUDING FOREGUT, MIDGUT, AND HINDGUT NEUROENDOCRINE TUMORS
- ODE-480 TO INCREASE THE NUMBER OF CIRCULATING MATURE NEUTROPHILS AND LYMPHOCYTES IN

PATENT AND EXCLUSIVITY TERMS**ORPHAN DRUG EXCLUSIVITY**

PATIENTS 12 YEARS OF AGE AND OLDER WITH WHIM SYNDROME (WARTS, HYPOGAMMAGLOBULINEMIA, INFECTIONS AND MYELOKATHEXIS)

- ODE-481 TREATMENT OF PEDIATRIC PATIENTS 2 YEARS OF AGE AND OLDER WITH ACTIVE POLYARTICULAR JUVENILE IDIOPATHIC ARTHRITIS WHO HAVE HAD AN INADEQUATE RESPONSE OR INTOLERANCE TO ONE OR MORE TUMOR NECROSIS FACTOR (TNF) BLOCKERS, AND FOR THE TREATMENT OF PEDIATRIC PATIENTS 2 YEARS OF AGE AND OLDER WITH ACTIVE PSORIATIC ARTHRITIS WHO HAVE HAD AN INADEQUATE RESPONSE OR INTOLERANCE TO ONE OR MORE TNF BLOCKERS
- ODE-482 TREATMENT OF ADULT PATIENTS WITH LOW- TO INTERMEDIATE-1 RISK MYELODYSPLASTIC SYNDROMES (MDS) WITH TRANSFUSION-DEPENDENT ANEMIA REQUIRING 4 OR MORE RED BLOOD CELL UNITS OVER 8 WEEKS WHO HAVE NOT RESPONDED TO OR HAVE LOST RESPONSE TO OR ARE INELIGIBLE FOR ERYTHROPOIESIS-STIMULATING AGENTS (ESA)
- ODE-483 TREATMENT OF ADULT AND PEDIATRIC PATIENTS 12 YEARS OF AGE AND OLDER WITH SOLID TUMORS THAT HAVE A NEUROTROPHIC TYROSINE RECEPTOR KINASE (NTRK) GENE FUSION, ARE LOCALLY ADVANCED OR METASTATIC OR WHERE SURGICAL RESECTION IS LIKELY TO RESULT IN SEVERE MORBIDITY, AND HAVE PROGRESSED FOLLOWING TREATMENT OR HAVE NO SATISFACTORY ALTERNATIVE THERAPY
- ODE-484 TREATMENT OF PEDIATRIC PATIENTS 2 YEARS OF AGE TO LESS THAN 12 YEARS OF AGE WITH ADVANCED OR METASTATIC MEDULLARY THYROID CANCER (MTC) WITH A RET MUTATION, AS DETECTED BY AN FDA-APPROVED TEST, WHO REQUIRE SYSTEMIC THERAPY
- ODE-485 TREATMENT OF PEDIATRIC PATIENTS 2 YEARS OF AGE TO LESS THAN 12 YEARS OF AGE WITH ADVANCED OR METASTATIC THYROID CANCER WITH A RET GENE FUSION, AS DETECTED BY AN FDA-APPROVED TEST, WHO REQUIRE SYSTEMIC THERAPY AND WHO ARE RADIOACTIVE IODINE-REFRACTORY (IF RADIOACTIVE IODINE IS APPROPRIATE)
- ODE-486 TREATMENT OF PRIMARY BILIARY CHOLANGITIS (PBC) IN ADULTS WHO HAVE HAD AN INADEQUATE RESPONSE TO URSODEOXYCHOLIC ACID (UDCA), OR IN PATIENTS UNABLE TO TOLERATE UDCA
- ODE-487 TREATMENT OF PEDIATRIC PATIENTS 2 YEARS OF AGE AND OLDER WITH LOCALLY ADVANCED OR METASTATIC SOLID TUMORS WITH A RET GENE FUSION, AS DETECTED BY AN FDA-APPROVED TEST, THAT HAVE PROGRESSED ON OR FOLLOWING PRIOR SYSTEMIC TREATMENT OR WHO HAVE NO SATISFACTORY ALTERNATIVE TREATMENT OPTIONS
- ODE-488 TREATMENT OF ADULT AND PEDIATRIC PATIENTS 12 YEARS OF AGE AND OLDER WITH SOLID TUMORS THAT HAVE A NEUROTROPIC TYROSINE RECEPTOR KINASE (NTRK) GENE FUSION, ARE LOCALLY ADVANCED OR METASTATIC OR WHERE SURGICAL RESECTION IS LIKELY TO RESULT IN SEVERE MORBIDITY, AND HAVE PROGRESSED FOLLOWING TREATMENT OR HAVE NO SATISFACTORY ALTERNATIVE THERAPY
- ODE-489 TREATMENT OF EXCESSIVE DAYTIME SLEEPINESS (EDS) IN PEDIATRIC PATIENTS 6 YEARS OF AGE AND OLDER WITH NARCOLEPSY
- ODE-490 TREATMENT OF CHOLESTATIC PRURITIS IN PATIENTS 12 MONTHS OF AGE TO LESS THAN 5 YEARS OF AGE WITH PROGRESSIVE FAMILIAL INTRAHEPATIC CHOLESTASIS (PFIC)
- ODE-491 TREATMENT OF ADULT AND PEDIATRIC PATIENTS 12 YEARS AND OLDER WITH GRADE 2 ASTROCYTOMA OR OLIGODENDROGLIOMA WITH A SUSCEPTIBLE ISOCITRATE DEHYDROGENASE-1 (IDH1) OR ISOCITRATE DEHYDROGENASE-2 (IDH2) MUTATION FOLLOWING SURGERY INCLUDING BIOPSY, SUB-TOTAL RESECTION, OR GROSS TOTAL RESECTION
- ODE-492 TREATMENT OF HYPOPARATHYROIDISM IN ADULTS
- ODE-493 TO SLOW KIDNEY FUNCTION DECLINE IN ADULTS WITH PRIMARY IMMUNOGLOBULIN A NEPHROPATHY (IGAN) WHO ARE AT RISK FOR DISEASE PROGRESSION, EXCLUDING THE USE PROVIDED FOR IN THE INDICATION APPROVED ON FEBRUARY 17, 2023
- ODE-494 TREATMENT OF CATAPLEXY OR EXCESSIVE DAYTIME SLEEPINESS (EDS) IN PEDIATRIC PATIENTS 7 YEARS OF AGE AND OLDER WITH NARCOLEPSY
- ODE-495 PROPHYLAXIS OF CYTOMEGALOVIRUS (CMV) DISEASE IN PEDIATRIC PATIENTS 12 YEARS OF AGE AND OLDER AND WEIGHING AT LEAST 40 KG WHO ARE KIDNEY TRANSPLANT RECIPIENTS AT HIGH RISK (DONOR CMV SEROPOSITIVE/RECIPIENT CMV SERONEGATIVE [D+/R-])
- ODE-496 TREATMENT OF NEUROLOGICAL MANIFESTATIONS OF NIEMANN-PICK DISEASE TYPE C (NPC) IN ADULT AND PEDIATRIC PATIENTS 2 YEARS OF AGE AND OLDER
- ODE-497 PROPHYLAXIS OF CYTOMEGALOVIRUS (CMV) INFECTION AND DISEASE IN PEDIATRIC PATIENTS 6 MONTHS OF AGE AND OLDER AND WEIGHING AT LEAST 6 KG WHO ARE CMV-SEROPOSITIVE RECIPIENTS [R+] OF AN ALLOGENEIC HEMATOPOIETIC STEM CELL TRANSPLANT (HSCT)
- ODE-498 TREATMENT OF NEUROLOGICAL MANIFESTATIONS OF NIEMANN-PICK DISEASE TYPE C (NPC) IN ADULTS AND PEDIATRIC PATIENTS WEIGHING GREATER THAN OR EQUAL TO 15 KG

PATENT AND EXCLUSIVITY TERMS

ADB 75 of 225

ORPHAN DRUG EXCLUSIVITY

ODE-499 TREATMENT OF ADULT PATIENTS WITH NEWLY DIAGNOSED PHILADELPHIA CHROMOSOME-POSITIVE CHRONIC MYELOID LEUKEMIA (PH+ CML) IN CHRONIC PHASE (CP), EXCLUDING PATIENTS WITH THE T315I MUTATION

ODE-500 TREATMENT OF ADULT PATIENTS WITH PREVIOUSLY TREATED PHILADELPHIA CHROMOSOME-POSITIVE CHRONIC MYELOID LEUKEMIA (PH+ CML) IN CHRONIC PHASE (CP), EXCLUDING PATIENTS PREVIOUSLY TREATED WITH TWO OR MORE TYROSINE KINASE INHIBITORS (TKIS), AND EXCLUDING PATIENTS WITH THE T315I MUTATION

ODE-501 ACUTE TREATMENT OF INTERMITTENT, STEREOTYPIC EPISODES OF FREQUENT SEIZURE ACTIVITY (I.E., SEIZURE CLUSTERS, ACUTE REPETITIVE SEIZURES) THAT ARE DISTINCT FROM A PATIENT'S USUAL SEIZURE PATTERN IN PATIENTS WITH EPILEPSY 2 TO 5 YEARS OF AGE

ODE-502 TREATMENT OF RELAPSED OR REFRACTORY MIXED-PHENOTYPE ACUTE LEUKEMIA WITH A LYSINE METHYLTRANSFERASE 2A GENE (KMT2A) TRANSLOCATION IN ADULT AND PEDIATRIC PATIENTS 1 YEAR AND OLDER.

ODE-503 ADJUNCTIVE TREATMENT TO GLUCOCORTICOID REPLACEMENT TO CONTROL ANDROGENS IN ADULTS AND PEDIATRIC PATIENTS 4 YEARS OF AGE AND OLDER WITH CLASSIC CONGENITAL ADRENAL HYPERPLASIA (CAH)

ODE-504 TREATMENT OF RELAPSED OR REFRACTORY ACUTE MYELOID LEUKEMIA WITH A LYSINE METHYLTRANSFERASE 2A GENE (KMT2A) TRANSLOCATION IN ADULT AND PEDIATRIC PATIENTS 1 YEAR AND OLDER

ODE-505 TREATMENT OF RELAPSED OR REFRACTORY ACUTE LYMPHOCYTIC LEUKEMIA WITH A LYSINE METHYLTRANSFERASE 2A GENE (KMT2A) TRANSLOCATION IN ADULT AND PEDIATRIC PATIENTS 1 YEAR AND OLDER

ODE-506 TREATMENT OF THE CARDIOMYOPATHY OF WILD-TYPE OR VARIANT TRANSTHYRETIN-MEDIATED AMYLOIDOSIS (ATTR-CM) IN ADULTS TO REDUCE CARDIOVASCULAR DEATH AND CARDIOVASCULAR-RELATED HOSPITALIZATION

PATENT USE

U-1 PREVENTION OF PREGNANCY

U-2 TREATMENT OR PROPHYLAXIS OF ANGINA PECTORIS AND ARRHYTHMIA

U-3 TREATMENT OF HYPERTENSION

U-4 PROVIDING PREVENTION AND TREATMENT OF EMESIS AND NAUSEA IN MAMMALS

U-5 METHOD OF PRODUCING BRONCHODILATION

U-6 METHOD OF PRODUCING SYMPATHOMIMETIC EFFECTS

U-7 INCREASING CARDIAC CONTRACTILITY

U-8 ACUTE MYOCARDIAL INFARCTION

U-9 CONTROL OF EMESIS ASSOCIATED WITH ANY CANCER CHEMOTHERAPY AGENT

U-10 DIAGNOSTIC METHOD FOR DISTINGUISHING BETWEEN HYPOTHALAMIC MALFUNCTIONS OR LESIONS IN HUMANS

U-11 TREATMENT OR PROPHYLAXIS OF CARDIAC DISORDERS

U-12 METHOD OF TREATING [A] HUMAN SUFFERING FROM DEPRESSION

U-13 A METHOD FOR TREATING ANXIETY IN A HUMAN SUBJECT IN NEED OF SUCH TREATMENT

U-14 ADJUNCTIVE THERAPY FOR THE PREVENTION AND TREATMENT OF HYPERAMMONEMIA IN THE CHRONIC MANAGEMENT OF PATIENTS WITH UREA CYCLE ENZYMOPATHIES

U-15 METHOD OF LOWERING INTRAOCULAR PRESSURE

U-16 USE IN LUNG SCANNING PROCEDURES

U-17 TREATMENT OF VENTRICULAR AND SUPRAVENTRICULAR ARRHYTHMIAS

U-18 METHOD FOR INHIBITING GASTRIC SECRETION IN MAMMALS

U-19 TREATMENT OF INFLAMMATION

U-20 A PROCESS FOR TREATING A PATIENT SUFFERING FROM PARKINSON'S SYNDROME AND IN NEED OF TREATMENT

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

- U-21 TREATMENT OF HUMANS SUFFERING UNDESIRE UROTOXIC SIDE EFFECTS CAUSED BY CYTOSTATICALLY ACTIVE ALKYLATING AGENTS
- U-22 METHOD OF COMBATTING PATHOLOGICALLY REDUCED CEREBRAL FUNCTIONS AND PERFORMANCE WEAKNESSES, CEREBRAL INSUFFICIENCY AND DISORDERS IN CEREBRAL CIRCULATION AND METABOLISM IN WARM-BLOODED ANIMALS
- U-23 METHOD FOR TREATING PROSTATIC CARCINOMA COMPRISING ADMINISTERING FLUTAMIDE
- U-24 METHOD FOR TREATING PROSTATE ADENOCARCINOMA COMPRISING ADMINISTERING AN ANTIANDROGEN INCLUDING FLUTAMIDE AND AN LHRH AGONIST
- U-25 REDUCING CHOLESTEROL IN CHOLELITHIASIS PATIENTS
- U-26 REDUCING CHOLESTEROL GALLSTONES AND/OR FRAGMENTS THEREOF
- U-27 DISSOLVING CHOLESTEROL GALLSTONES AND/OR FRAGMENTS THEREOF
- U-28 CEREBRAL, CORONARY, PERIPHERAL, VISCERAL AND RENAL ARTERIOGRAPHY, AORTOGRAPHY AND LEFT VENTRICULOGRAPHY
- U-29 CT IMAGING OF THE HEAD AND BODY, AND INTRAVENOUS EXCRETORY UROGRAPHY
- U-30 CEREBRAL ANGIOGRAPHY, AND VENOGRAPHY
- U-31 INTRA-ARTERIAL DIGITAL SUBTRACTION ANGIOGRAPHY
- U-32 PALLIATIVE TREATMENT OF PATIENTS WITH OVARIAN CARCINOMA RECURRENT AFTER PRIOR CHEMOTHERAPY, INCLUDING PATIENTS WHO HAVE BEEN PREVIOUSLY TREATED WITH CISPLATIN
- U-33 TREATING VIRAL INFECTIONS IN A MAMMAL
- U-34 TREATING VIRAL INFECTIONS IN A WARM-BLOODED ANIMAL
- U-35 TREATING CYTOMEGALOVIRUS IN A HUMAN WITH AN INJECTABLE COMPOSITION
- U-36 METHODS OF TREATING BACTERIAL ILLNESSES
- U-37 METHOD OF TREATING GASTROINTESTINAL DISEASE
- U-38 TREATMENT OF PAROXYSMAL SUPRAVENTRICULAR TACHYCARDIA
- U-39 ANGINA PECTORIS
- U-40 METHOD OF TREATMENT OF BURNS
- U-41 METHOD OF TREATING CARDIAC ARRHYTHMIAS
- U-42 ADJUVANT TREATMENT IN COMBINATION WITH FLUOROURACIL AFTER SURGICAL RESECTION IN PATIENTS WITH DUKES' STAGE C COLON CANCER
- U-43 MANAGEMENT OF CHRONIC PAIN IN PATIENTS REQUIRING OPIOID ANALGESIA
- U-44 RELIEF OF NAUSEA AND VOMITING
- U-45 TREATMENT OF INFLAMMATION AND ANALGESIA
- U-46 TREATMENT OF PANIC DISORDER
- U-47 STIMULATION OF THE RELEASE OF GROWTH HORMONE
- U-48 ANALGESIA
- U-49 SYMPTOMATIC CANCER-RELATED HYPERCALCEMIA
- U-50 USE IN TREATING INFLAMMATORY DERMATOSES
- U-51 BLOOD POOL IMAGING, INCLUDING CARDIAC FIRST PASS AND GATED EQUILIBRIUM IMAGING AND FOR DETECTION OF SITES OF GASTROINTESTINAL BLEEDING
- U-52 TREATMENT OF ADULT AND PEDIATRIC PATIENTS(OVER SIX MONTHS OF AGE) WITH ADVANCED HIV INFECTION
- U-53 HYPERCALCEMIA OF MALIGNANCY
- U-54 REVERSAL AGENT OR ANTAGONIST OF NONDEPOLARIZING NEUROMUSCULAR BLOCKING AGENTS
- U-55 TREATMENT OF PAIN
- U-56 AID TO SMOKING CESSATION
- U-57 OPHTHALMIC USE OF NORFLOXACIN
- U-58 METHOD OF TREATING INFLAMMATORY INTESTINAL DISEASES
- U-59 METHOD OF TREATING HYPERCHOLESTEROLEMIA
- U-60 NASAL ADMINISTRATION OF BUTORPHANOL

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-61 CEREBRAL AND PERIPHERAL ARTERIOGRAPHY AND CT IMAGING OF THE HEAD

U-62 CORONARY ARTERIOGRAPHY, LEFT VENTRICULOGRAPHY, CT IMAGING OF THE BODY, INTRAVENOUS EXCRETORY UROGRAPHY, INTRAVENOUS DIGITAL SUBTRACTION ANGIOGRAPHY AND VENOGRAPHY

U-63 ISOPRENALINE ANTAGONISM ON THE HEART RATE OR BLOOD PRESSURE

U-64 TREATMENT OF VIRAL INFECTIONS

U-65 METHOD OF TREATMENT OF A PATIENT INFECTED WITH HIV

U-66 TRIPHASIC REGIMEN

U-67 METHOD OF INDUCING ANESTHESIA IN A WARM BLOODED ANIMAL

U-68 TREATMENT OF ACTINIC KERATOSIS

U-69 TREATMENT OF PNEUMOCYSTIS CARINII INFECTIONS

U-70 TREATMENT OF TRANSIENT INSOMNIA

U-71 METHOD OF TREATMENT OF HEART FAILURE

U-72 TREATMENT OF MIGRAINE

U-73 METHOD OF TREATING DISEASES OR INFECTIONS CAUSED BY MYCETES

U-74 METHOD OF PROVIDING HYPNOTIC EFFECT

U-75 RELIEF OF OCULAR ITCHING DUE TO SEASONAL ALLERGIC CONJUNCTIVITIS

U-76 USE TO IMAGE A SUBJECT WITH A MAGNETIC RESONANCE IMAGING SYSTEM

U-77 TREATMENT OF SYMPTOMS OF SEASONAL ALLERGIC RHINITIS

U-78 ULCERATIVE COLITIS

U-79 SYMPTOMATIC TREATMENT OF PATIENTS WITH NOCTURNAL HEARTBURN DUE TO GERD

U-80 METHOD OF TREATING OCULAR BACTERIAL INFECTIONS

U-81 RELIEF OF SYMPTOMS ASSOCIATED WITH SEASONAL ALLERGIC RHINITIS

U-82 TREATMENT FOR DEMENTIA IN PATIENTS WITH ALZHEIMER'S DISEASE

U-83 TREATMENT OF SEIZURES

U-84 A METHOD OF BLOCKING THE UPTAKE OF MONOAMINES BY BRAIN NEURONS IN ANIMALS

U-85 NASAL TREATMENT OF SEASONAL AND PERENNIAL ALLERGIC RHINITIS SYMPTOMS

U-86 METHOD OF TREATING CERTAIN FORMS OF EPILEPSY

U-87 METHOD FOR NONINVASIVE ADMINISTRATION OF SEDATIVES, ANALGESICS, AND ANESTHETICS

U-88 TREATMENT OF MODERATE PLAQUE PSORIASIS

U-89 TREATMENT OR PROPHYLAXIS OF EMESIS

U-90 TREATMENT OF PSYCHOTIC DISORDERS

U-91 ALTERNATIVE THERAPY TO TRIMETHOPRIM-SULFAMETHOXAZOLE FOR TREATMENT OF MODERATE-TO-SEVERE PNEUMOCYSTIS CARINII PNEUMONIA IN IMMUNOCOMPROMISED AND AIDS PATIENTS

U-92 TREATMENT OF DIABETIC NEPHROPATHY IN PATIENTS WITH TYPE I INSULIN DEPENDENT DIABETES MELLITUS AND RETINOPATHY

U-93 USE AS AN ANTIHISTAMINE/DECONGESTANT

U-94 TREATMENT-ADULTS W/ ADVANCED HIV, INTOLERANT OF APPROVED THERAPIES, INTOLERANT OF APPROVED THERAPIES W/PROVEN BENEFIT OR HAVE EXPERIENCED CLINICAL/IMMUNOLOGICAL DETERIORATION WHILE RECEIVING..OR FOR WHOM SUCH THERAPIES-CONTRAINDICATED

U-95 SHORT TERM MANAGEMENT OF MODERATE PRURITIS IN ADULTS WITH ATOPIC DERMATITIS AND LICHEN SIMPLEX CHRONICUS

U-96 METHOD OF TREATING VARICELLA ZOSTER (SHINGLES) INFECTIONS

U-97 A METHOD OF TREATING A PATIENT IN NEED OF MEMORY ENHANCEMENT

U-98 A METHOD OF INDUCING REGRESSION OF LEUKEMIA CELL GROWTH IN A MAMMAL

U-99 METHOD OF PROVIDING POTASSIUM TO A SUBJECT IN NEED OF POTASSIUM

U-100 METHOD OF TREATING OCULAR INFLAMMATION

U-101 ADJUNCT TO CONVENTIONAL CT OR MRI IMAGING IN THE LOCALIZATION OF STROKE IN PATIENTS IN WHOM STROKE HAS ALREADY BEEN DIAGNOSED

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-102 METHOD OF HORMONALLY TREATING MENOPAUSAL OR POST-MENOPAUSAL DISORDERS IN WOMEN

U-103 TREATMENT OF OCULAR HYPERTENSION

U-104 TREATMENT OF AQUEOUS HUMOR FORMATION AND INTRAOCULAR PRESSURE

U-105 EMESIS

U-106 TREATMENT OF EPILEPSY

U-107 TREATMENT OF HYPERTENSION AND ANGINA PECTORIS

U-108 SHORT-TERM TREATMENT OF ACTIVE DUODENAL ULCER, GASTROESOPHAGEAL REFLUX DISEASE (GERD), SEVERE EROSIIVE ESOPHAGITIS, POORLY RESPONSIVE SYMPTOMATIC GERD AND PATHOLOGIAL HYPERSECRETORY CONDITIONS AND MAINTENANCE HEALING OF EROSIIVE ESOPHAGITIS

U-109 ADJUNCT DIET IN THE TX OF ELEVATED TOTAL CHOLESTEROL AND LDL-C LEVELS IN PTS W/PRIMARY HYPERCHOLESTEROLEMIA WHOSE RESPONSE TO DIETARY RESTRICTION OF SAT FAT AND CHOLESTEROL AND OTHER NONPHARMACOLOGICAL MEASURES HAS NOT BEEN ADEQUATE

U-110 USE AS A RETRIEVABLE PESSARY

U-111 DIABETES

U-112 CONTRACEPTION

U-113 METHOD OF CONDUCTING RADIOLOGICAL EXAMINATION OF A PATIENT BY ADMINISTERING TO SAID PATIENT A RADIOPAQUE AMOUNT OF IOPROMIDE

U-114 USE FOR INHIBITING BONE RESORPTION

U-115 USE OF VASODILATORS TO EFFECT AND ENHANCE AN ERECTION (AND THUS TREAT ERECTILE DYSFUNCTION), BY INJECTION INTO THE PENIS

U-116 METHOD OF MYOCARDIAL IMAGING

U-117 TREATMENT OF OCULAR ALLERGIC RESPONSE IN HUMAN EYES

U-118 METHOD OF LOWERING BLOOD SUGAR LEVEL

U-119 TREATMENT OF NASAL HYPERSECRETION

U-120 CONTROLLING OR PREVENTING POST-OPERATIVE INTRAOCULAR PRESSURE RISES ASSOCIATED WITH OPHTHALMIC LASER SURGICAL PROCEDURES

U-121 METHOD OF TREATING CONDITIONS MEDIATED THROUGH HISTAMINE H2-RECEPTORS

U-122 A THERAPEUTIC METHOD FOR CONTROLLING THROMBOSIS

U-123 METHOD FOR CONTROLLING THROMBOSIS AND DECREASING BLOOD HYPERCOAGULATION AND HEMORRHAGING RISKS

U-124 TREATMENT OF ACNE

U-125 TREATMENT NEUROGENERATIVE DISEASES

U-126 TREATMENT OF GASTRITIS

U-127 METHOD OF PRODUCING NEUROMUSCULAR BLOCKADE

U-128 METHOD FOR TREATMENT OF TUMORS

U-129 METHOD TO DESTROY OR IMPAIR TARGET CELLS

U-130 MANAGEMENT OF PATIENTS WITH MASTOCYTOSIS

U-131 PHOTODAMAGED SKIN

U-132 INHIBITING HIV PROTEASE

U-133 MANAGEMENT OF OBESITY INCLUDING WEIGHT LOSS AND MAINTENANCE IN PATIENTS ON A REDUCED-CALORIE DIET

U-134 TREATMENT OF ACNE VULGARIS

U-135 ANTITUMOR AGENT

U-136 PROCESS FOR WASTE NITROGEN REMOVAL

U-137 METHOD OF TREATING BACTERIAL VAGINOSIS

U-138 TREATMENT OF ALLERGIC RHINITIS

U-139 TREATMENT OF ALLERGIC REACTIONS

U-140 USE OF NORVIR TO INHIBIT HIV PROTEASE OR TO INHIBIT AN HIV INFECTION

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-141 TREATMENT OF ULCERATIVE COLITIS

U-142 METHOD OF TREATING ALLERGIC REACTIONS IN A MAMMAL BY USING THIS ACTIVE METABOLITE

U-143 BIODEGRADABLE SUPERPARAMAGNETIC METAL OXIDES AS CONTRAST AGENTS FOR MR IMAGING

U-144 BIOLOGICALLY DEGRADABLE SUPERPARAMAGNETIC MATERIALS FOR USE IN CLINICAL APPLICATIONS

U-145 BIOLOGICALLY DEGRADABLE SUPERPARAMAGNETIC PARTICLES FOR USE AS NUCLEAR MAGNETIC RESONANCE IMAGING AGENTS

U-146 METHOD OF TREATING SUSCEPTIBLE NEOPLASMS IN MAMMALS

U-147 DETECTION OF GASTROINTESTINAL DISORDERS AND THE SUBSEQUENT BREATH COLLECTION AND MEASUREMENT OF $^{13}\text{CO}_2$

U-148 DEVICE FOR COLLECTING A BREATH SAMPLE

U-149 METHOD OF TREATING AN ANIMAL, INCLUDING A HUMAN SUFFERING FROM OR SUSCEPTIBLE TO PSYCHOSIS, ACUTE MANIA OR MILD ANXIETY STATES

U-150 METHOD OF USE FOR CONTROLLING HYPERGLYCEMIA BY ADMINISTRATION OF THIS SUSTAINED RELEASE DOSAGE FORM OF GLIPIZIDE

U-151 RELIEF OF SYMPTOMS OF THE COMMON COLD

U-152 METHOD OF TREATING ANXIETY RELATED DISORDERS INCLUDING OBSESSIVE COMPULSIVE DISORDER

U-153 TREATMENT OF INITIAL EPISODE GENITAL HERPES

U-154 METHOD OF TREATING ANIMALS SUFFERING FROM AN APPETITE DISORDER

U-155 TREATMENT OF ERECTILE DYSFUNCTION

U-156 METHOD OF PROVIDING ANESTHESIA

U-157 TREATMENT OF A HUMAN SUFFERING FROM VITAMIN B12 DEFICIENCY

U-158 ANGINA

U-159 TREATMENT OF INTERSTITIAL CYSTITIS

U-160 TREATMENT OF BACTERIAL INFECTIOUS DISEASE

U-161 METHOD OF INHIBITING CHOLESTEROL BIOSYNTHESIS IN A PATIENT

U-162 METHOD OF USE TO INHIBIT CHOLESTEROL SYNTHESIS IN A HUMAN SUFFERING FROM HYPERCHOLESTEROLEMIA

U-163 METHOD OF USING TROGLITAZONE TO TREAT IMPAIRED GLUCOSE TOLERANCE TO PREVENT OR DELAY THE ONSET OF NONINSULIN-DEPENDENT DIABETES MELLITUS

U-164 METHOD OF USING TROGLITAZONE TO PREVENT OR DELAY THE ONSET OF NONINSULIN-DEPENDENT DIABETES MELLITUS IN A DEFINED POPULATION OF PATIENTS

U-165 TREATMENT OF SYMPTOMATIC BENIGN PROSTATIC HYPERPLASIA

U-166 TREATMENT OF H.PYLORI-ASSOCIATED DUODENAL ULCER

U-167 METHOD FOR TREATING HIV-1 INFECTION

U-168 METHOD OF INHIBITING LIPOXYGENASE ACTIVITY IN A MAMMAL WHICH IS THE MODE OF ACTION IN THE TREATMENT OF ASTHMA

U-169 METHODS OF USING THE COMPOUND/DRUG PRODUCT AS A CONTRAST AGENT IN MAGNETIC RESONANCE IMAGING

U-170 METHOD OF OBTAINING AN MR IMAGE USING THE COMPOSITION/DRUG PRODUCT AS A CONTRAST AGENT

U-171 METHODS OF USING THE COMPOUND/DRUG PRODUCT AS AN ORAL CONTRAST AGENT IN MAGNETIC RESONANCE IMAGING OF THE GASTROINTESTINAL TRACT

U-172 TREATMENT OF GENITAL WARTS

U-173 ADMINISTRATION TO A HOST SUFFERING FROM GESTATIONAL DIABETES

U-174 USE AS AN ANTIHISTAMINE AGENT

U-175 METHOD OF TREATING MALIGNANT TUMORS

U-176 METHOD OF TREATING A PATIENT SUFFERING FROM LISTED CONDITIONS, INCLUDING SPECIFIC PSYCHOSES

U-177 FUNGICIDE

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-178 FACILITATED ADHERENCE OF AGENTS TO SKIN

U-179 ENHANCED CUTANEOUS PENETRATION OF A DERMALLY-APPLIED PHARMACOLOGICALLY ACTIVE AGENT

U-180 TREATMENT OF ADULT AND PEDIATRIC PATIENTS (OVER 6 MONTHS OF AGE) WITH ADVANCED HIV INFECTION

U-181 PRODUCING ALPHA ADRENERGIC ANTAGONISTIC ACTION IN A HOST

U-182 USE OF SALMETEROL IN PATIENTS WITH REVERSIBLE AIRWAY OBSTRUCTION

U-183 TREATMENT OF CONDITIONS CAUSED BY DISTURBANCE OF NEURONAL 5HT FUNCTION

U-184 TREATING ALLERGIC EYE DISEASES IN HUMANS

U-185 METHOD OF TREATING HYPERTENSION

U-186 METHOD FOR TREATING GI DISORDERS CAUSED BY H. PYLORI WHICH COMPRISES ADMINISTRATION OF RANITIDINE BISMUTH CITRATE AND CLARITHROMYCIN FOR A GREATER THAN ADDITIVE EFFECT

U-187 THERAPEUTIC TREATMENT OF CALCIFIC TUMORS

U-188 TREATMENT OF H.PYLORI ASSOCIATED DUODENAL ULCER

U-189 ENHANCEMENT OF THE BIOAVAILABILITY OF THE DRUG SUBSTANCE

U-190 USE OF RITONAVIR IN COMBINATION WITH ANY REVERSE TRANSCRIPTASE INHIBITOR

U-191 METHOD OF TREATMENT FOR CONTROLLING AND LOWERING INTRAOCULAR PRESSURE IN A HUMAN

U-192 USE IN TREATING ALLERGIC REACTIONS

U-193 PSORIASIS

U-194 TREATING ANGINA PECTORIS AND HIGH BLOOD PRESSURE

U-195 METHOD FOR THE DIAGNOSIS OF GASTROINTESTINAL DISORDERS BY UREA ISOTOPE OR NITROGEN LABELED CARBON

U-196 TREATMENT OF METASTATIC BREAST CANCER IN POSTMENOPAUSAL WOMEN WITH ESTROGEN RECEPTOR POSITIVE TUMORS

U-197 USE IN COMBINATION WITH CERTAIN LHRH ANALOGUES FOR THE TREATMENT OF ADVANCED PROSTATE CANCER

U-198 TREATMENT METASTATIC CARCINOMA OF OVARY AFTER 1ST LINE FAILURE OR SUBSEQUENT CHEMOTHERAPY, TREATMENT OF BREAST CANCER AFTER FAILURE OF COMBINATION CHEMOTHERAPY FOR METASTATIC DISEASE AND 2ND LINE TREATMENT OF AIDS RELATED KAPOSI'S SARCOMA

U-199 METHOD OF TREATING INFECTIOUS UPPER GI TRACT DISORDERS CAUSED BY CAMPYLOBACTER PYLORIDIS INFECTION COMPRISING ADMINISTRATION OF A BISMUTH AGENT AND AN ANTIMICROBIAL AGENT

U-200 METHOD OF TREATING GI DISORDERS COMPRISING ADMINISTRATION OF A BISMUTH-CONTAINING AGENT AND H2 RECEPTOR BLOCKING ANTI-SECRETORY AGENT

U-201 METHOD OF TREATING GI DISORDERS COMPRISING ADMINISTRATION OF CAMPYLOBACTER-INHIBITING ANTIMICROBIAL AGENT AND H2 RECEPTOR BLOCKING ANTI-SECRETORY AGENT

U-202 METHOD OF TREATING PEPTIC ULCER DISEASE CAUSED BY CAMPYLOBACTER PYLORIDIS COMPRISING ORAL ADMINISTRATION OF 50 TO 5,000MG BISMUTH DAILY FOR 3-56 DAYS

U-203 TREATMENT OF ADVANCED BREAST CANCER IN POSTMENOPAUSAL WOMEN WITH DISEASE PROGRESSION FOLLOWING ANTIESTROGEN THERAPY

U-204 USE OF TAXOL IN COMBINATION WITH G-CSF FOR TREATMENT OF PATIENTS WITH AIDS-RELATED KAPOSI'S SARCOMA

U-205 METHOD FOR TREATING HEARTBURN

U-206 METHOD OF USING FSH ALONE, WITHOUT THE PRESENCE OF EXOGENEOUS LH, IN IN VITRO FERTILIZATION

U-207 USE AS NASAL SPRAY

U-208 VAGINAL ADMINISTRATION USING SPECIFIED FORMULATION

U-209 VAGINAL ADMINISTRATION OF PROGESTERONE USING SPECIFIED FORMULATION

U-210 METHOD OF TREATING CONGESTIVE HEART FAILURE

U-211 USE IN PATIENTS WITH REVERSIBLE AIRWAY OBSTRUCTION

U-212 METHOD OF TREATMENT OF PARKINSON'S DISEASE

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-213 METHOD OF INHIBITING CHOLESTEROL BIOSYNTHESIS AND TREATING HYPERCHOLESTEROLEMIA AND METHOD FOR TREATING HYPERLIPIDEMIA

U-214 USE AS A BLOOD GLUCOSE-LOWERING AGENT

U-215 TREATMENT OF EPILEPSY TWICE DAILY. TREATING A PATIENT BY ADMINISTERING CARBAMAZEPINE IN A DOSAGE FORM CAPABLE OF MAINTAINING BLOOD CONCENTRATION FROM 4-12MCG/ML OVER 12 HOURS

U-216 TREATMENT OF ADENOCARCINOMA, INCLUDING STAGE B2-C BY ADMINISTERING AN AGONIST OF LH-RH AND FLUTAMIDE

U-217 METHOD OF PRODUCING ANESTHESIA

U-218 METHOD FOR LIMITING THE POTENTIAL FOR MICROBIAL GROWTH IN THE DRUG PRODUCT

U-219 TREATMENT OF PARKINSON'S DISEASE

U-220 METHOD OF DIAGNOSIS

U-221 SELECTIVE VASODILATION BY CONTINUOUS ADENOSINE INFUSION

U-222 METHOD OF TREATING PAGET'S DISEASE USING ACTONEL

U-223 TREATMENT OF BACTERIAL CONJUNCTIVITIS CAUSED BY SUSCEPTIBLE STRAINS OF MICROORGANISMS

U-224 CONTROLLING INTRAOCULAR PRESSURE

U-225 METHOD FOR DELIVERY

U-226 METHOD OF ENHANCING THE DISSOLUTION PROFILE OF A PHARMACEUTICAL FROM A SOLID DOSAGE FORM CONTAINING THE PHARMACEUTICAL AND SIMETHICONE

U-227 NASAL ADMINISTRATION

U-228 ASTHMA

U-229 CARDIAC INSUFFICIENCY (CONGESTIVE HEART FAILURE)

U-230 PREVENTION OF ACUTE CARDIAC ISCHEMIC EVENTS

U-231 USE IN PARKINSON'S DISEASE

U-232 METHOD OF TREATING MIGRAINE

U-233 DECREASING MORTALITY CAUSED BY CONGESTIVE HEART FAILURE

U-234 METHOD OF USING RIBAVIRIN TO TREAT VIRAL INFECTIONS IN MAMMALS

U-235 METHOD OF MODULATING TH1 AND TH2 RESPONSE IN ACTIVATED T CELLS OF A HUMAN COMPRISING ADMINISTERING RIBAVIRIN TO THE T CELLS IN A DOSAGE WHICH PROMOTES THE TH1 RESPONSE AND SUPPRESSES THE TH2 RESPONSE

U-236 TREATING MALE PATTERN BALDNESS WITH 0.05 TO 3.0MG/DAY

U-237 METHOD OF PERFORMING NMR IMAGING WITH A PATIENT COMPRISING ADMINISTERING TO THE PATIENT AN EFFECTIVE AMOUNT OF CONTRAST AGENT DISCLOSED IN THE CLAIMS

U-238 IMAGING A BODY TISSUE AND SUBJECTING TO NMR TOMOGRAPHY, ADMINISTERING AN AMOUNT OF PHARMACEUTICAL AGENT FOR AFFECTING THE RELAXATION TIMES OF ATOMS IN BODY TISSUES UNDERGOING NMR DIAGNOSIS, WHEREBY THE IMAGE CONTRAST IS ENHANCED....

U-239 TREATING OR CONTROLLING OCULAR INFLAMMATION WHICH COMPRISES TOPICALLY ADMINISTERING TO AFFECTED EYE A COMPOSITION COMPRISING AN NSAID, A POLYMERIC QUATERNARY AMMONIUM COMPOUND AND BORIC ACID

U-240 TREATMENT OF ACUTE MIGRAINE ATTACKS

U-241 FOR SHORT-TERM TREATMENT ACTIVE DUODENAL ULCER, MAINTENANCE THERAPY FOR DUODENAL ULCER PATIENTS AT REDUCED DOSAGE AFTER HEALING OF ACTIVE ULCER, SHORT-TERM TREATMENT ACTIVE BENIGN GASTRIC ULCER & GERD, PATHOLOGICAL HYPERSECRETORY CONDITIONS

U-242 USE OF FOLLITROPIN ALPHA ALONE IN IN-VITRO FERTILIZATION

U-243 TOPICAL ADMINISTRATION

U-244 PLATELET AGGREGATION INHIBITORS

U-245 TREATMENT OF SEBORRHEA DERMATITIS IN HUMANS

U-246 PHOSPHATE BINDING

U-247 TREATMENT OF RHEUMATOID ARTHRITIS

U-248 TREATMENT OF HIV

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-249 METHOD OF TREATING ALLERGIC OR NON-ALLERGIC RHINITIS IN PATIENTS BY ADMINISTERING AEROSOLIZED PARTICLES OF MOMETASONE FUROATE

U-250 TREATMENT OF HEPATITIS B INFECTION

U-251 USE OF TROGLITAZONE IN COMBINATION WITH SULFONYLUREAS IN THE TREATMENT OF TYPE II DIABETES

U-252 METHOD OF TREATING A HUMAN SUBJECT HAVING GAUCHER'S DISEASE

U-253 ORAL TRANSMUCOSAL USE

U-254 USE OF AGGRASTAT IN COMBINATION WITH HEPARIN

U-255 IMPROVED WAKEFULNESS IN PATIENTS WITH EXCESSIVE DAYTIME SLEEPINESS ASSOCIATED WITH NARCOLEPSY

U-256 TREATMENT OF HIV INFECTION IN COMBINATION WITH ONE OR MORE ADDITIONAL HIV ANTIVIRAL AGENTS

U-257 TREATMENT OF HIV INFECTION

U-258 TREATMENT OF NEURODEGENERATIVE DISEASES

U-259 TREATMENT OF ANDROGENIC ALOPECIA BY ORAL ADMINISTRATION DRUG SUBSTANCE

U-260 REDUCTION OF INTRAOCULAR PRESSURE IN PATIENTS WITH OPEN ANGLE GLAUCOMA AND OCULAR HYPERTENSION WHO ARE INTOLERANT OF OTHER IOP LOWERING MEDICATIONS OR INSUFFICIENTLY RESPONSIVE TO ANOTHER IOP LOWERING MEDICATION

U-261 TREATING BENIGN PROSTATIC HYPERPLASIA WITH A GENUS OF COMPOUNDS, INCLUDING FINASTERIDE

U-262 TREATING BENIGN PROSTATIC HYPERTROPHY WITH FINASTERIDE

U-263 METHOD OF TREATING A MALIGNANT CONDITION THROUGH INTRAVASCULAR ADMINISTRATION OF BUSULFAN. METHOD FOR TREATING LEUKEMIA OR LYMPHOMA IN A PATIENT UNDERGOING A BONE MARROW TRANSPLANT THROUGH INTRAVENOUS ADMINISTRATION OF BUSULFAN

U-264 METHOD OF TREATING A MALIGNANT DISEASE THROUGH PARENTERAL ADMINISTRATION OF BUSULFAN. METHOD FOR TREATING A PATIENT UNDERGOING A BONE MARROW TRANSPLANT THROUGH INTRAVASCULAR ADMINISTRATION OF BUSULFAN

U-265 USE AS LAXATIVE

U-266 RELIEF OF THE SIGNS AND SYMPTOMS OF OSTEOARTHRITIS; RELIEF OF THE SIGNS AND SYMPTOMS OF RHEUMATOID ARTHRITIS IN ADULTS; MANAGEMENT OF ACUTE PAIN IN ADULTS; TREATMENT OF PRIMARY DYSMENORRHEA; ACUTE TREATMENT OF MIGRAINE ATTACKS IN ADULTS

U-267 PREVENTING HEARTBURN EPISODES FOLLOWING INGESTION OF HEARTBURN-INDUCING FOOD/BEVERAGE, COMPRISING ADMIN TO PT, 30 MIN PRIOR TO CONSUMPTION BY THE PT THE FOOD/BEVERAGE, A COMPOSITION COMPRISING 10MG FAMOTIDINE

U-268 ACROMEGALY

U-269 EXCESS GH-SECRETION OR GASTRO-INTESTINAL DISORDERS

U-270 METHOD OF IMPROVING THE TIME FOR ADMINISTRATION OR THE TIME BETWEEN CHANGES OF GIVING SETS FOR THE DRUG PRODUCT

U-271 METHOD OF TREATING TUMORS

U-272 METHOD OF TREATING CARCINOMA

U-273 CUTANEOUS T-CELL LYMPHOMA

U-274 ZANAMIVIR FOR INHALATION

U-275 METHOD OF USE OF THE DRUG SUBSTANCE

U-276 METHOD OF USE OF LEVOBUPIVACAINE

U-277 NEUROLOGICAL AND OTHER DISORDERS (TREATMENT OF EPILEPSY, BID ORAL DOSING)

U-278 METHOD OF USE OF THE INDICATION OF THE DRUG PRODUCT

U-279 METHOD OF USE OF THE APPROVED PRODUCT

U-280 TREATING PRECIPITATED ACUTE URINARY RETENTION WITH FINASTERIDE

U-281 ANTIMYCOTIC USES, SPECIFICALLY TREATMENT OF ONYCHOMYCOSIS

U-282 METHOD OF TREATING BACTERIAL INFECTIONS

U-283 METHOD FOR TREATING MENOPAUSAL SYMPTOMS IN A POSTMENOPAUSAL FEMALE

U-284 MENOPAUSAL AND POSTMENOPAUSAL DISORDERS (INCLUDING VASOMOTOR SYMPTOMS ASSOCIATED

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

WITH MENOPAUSE, AND VULVAR AND VAGINAL ATROPHY) AND OSTEOPOROSIS

U-285 DEPRESSION AND SOCIAL ANXIETY DISORDER/SOCIAL PHOBIA

U-286 DEPRESSION

U-287 TREATMENT OR PREVENTION OF OSTEOPOROSIS

U-288 THERAPY OF INFLUENZA

U-289 TREATMENT OF NON-HYPERKERATOTIC ACTINIC KERATOSES OF FACE AND SCALP

U-290 INHIBITING TRANSPLANT REJECTION USING RAPAMYCIN (SIROLIMUS)

U-291 INHIBITING TRANSPLANT REJECTION USING RAPAMYCIN (SIROLIMUS) IN COMBINATION WITH CYCLOSPORIN

U-292 INHIBITING TRANSPLANT REJECTION USING RAPAMYCIN (SIROLIMUS) IN COMBINATION WITH AZATHIOPRINE

U-293 INHIBITING TRANSPLANT REJECTION USING RAPAMYCIN (SIROLIMUS) IN COMBINATION WITH A CORTICOSTEROID

U-294 TREATMENT OF HYPERPIGMENTARY DISORDERS

U-295 TREATMENT OF SEASONAL AND PERENNIAL ALLERGIC RHINITIS SYMPTOMS

U-296 TREATING MIGRAINE PAIN AND ONE OR MORE OF A CLUSTER OF SYMPTOMS CHARACTERISTIC OF A MIGRAINE ATTACK SYMPTOMS BEING SELECTED FROM PHOTOPHOBIA, PHONOPHOBIA NAUSEA AND FUNCTIONAL DISABILITY

U-297 PREVENTION OR TREATMENT OF REVERSIBLE VASOCONSTRICTION BY THE INHALATION OF NITRIC OXIDE WITH AN OXYGEN CONTAINING GAS

U-298 METHOD OF COMBATING BACTERIA IN A PATIENT

U-299 TREATMENT OF ADENOMATOUS POLYPS

U-300 INDICATED FOR THE REDUCTION OF ELEVATED TOTAL AND LDL CHOLESTEROL LEVELS IN PATIENTS WITH PRIMARY HYPERCHOLESTEROLEMIA

U-301 USE OF TROGLITAZONE IN COMBINATION WITH SULFONYLUREAS AND BIGUANIDES IN THE TREATMENT OF TYPE II DIABETES

U-302 TO REDUCE THE RISK OF STROKE IN PATIENTS WHO HAVE HAD TRANSIENT ISCHEMIA OF THE BRAIN OR COMPLETED ISCHEMIC STROKE DUE TO THROMBOSIS

U-303 METHOD OF USE PATENT-PRODUCT APPROVED FOR TREATMENT OF OSTEOPOROSIS, PAGET'S DISEASE, PREVENTION AND TREATMENT OF GLUCOCORTICOID INDUCED OSTEOPOROSIS

U-304 A METHOD OF TREATMENT OF A CONDITION INVOLVING AN ANTIBODY ANTIGEN REACTION

U-305 METHODS FOR USING THE DRUG PRODUCT

U-306 TREATMENT OF POST-MENOPAUSAL UROGENITAL SYMPTOMS ASSOCIATED WITH ESTROGEN DEFICIENCY

U-307 CLAIMS AN OLANZAPINE POLYMORPH USEFUL FOR TREATING ANY NUMBER OF LISTED CONDITIONS, INCLUDING SPECIFIC PSYCHOSES, EMPLOYING OLANZAPINE AS PER THE INDICATION OF THIS NDA

U-308 CLAIMS A SOLID ORAL FORMULATION INCLUDING TABLETS AND GRANULES OF OLANZAPINE USEFUL FOR TREATING ANY NUMBER OF LISTED CONDITIONS, INCLUDING SPECIFIC PSYCHOSES, EMPLOYING OLANZAPINE AS PER THE INDICATIONS OF THIS NDA

U-309 TREATING SJOEGREN SYNDROME

U-310 TREATMENT OF XEROSTOMIA

U-311 HORMONE REPLACEMENT

U-312 PANIC DISORDER, OBSESSIVE-COMPULSIVE DISORDER, POSTTRAUMATIC STRESS DISORDER

U-313 TREATMENT OF CONGESTIVE HEART FAILURE

U-314 METHOD FOR TREATING HYPERPARATHYROIDISM WHICH COMPRISES SUPPRESSING PARATHYROID ACTIVITY

U-315 METHOD FOR ADMINISTERING DRUG TO GASTROINTESTINAL TRACT

U-316 METHOD OF TREATING A SUBJECT SUFFERING FROM PROSTATE CANCER

U-317 METHOD OF USING TROGLITAZONE TO TREAT PATIENTS HAVING INSULIN RESISTANCE

U-318 TREATMENT OF PATIENTS WITH AN OVERACTIVE BLADDER WITH SYMPTOMS OF URINARY FREQUENCY, URGENCY, OR URGE INCONTINENCE

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-319 TREATMENT OF MICROBIAL INFECTIONS

U-320 INHIBITING OR ELIMINATING ACUTE MYELOID LEUKEMIA

U-321 REDUCTION OF ELEVATED IPTH LEVELS IN THE MGT OF SECONDARY HYPERPARATHYROIDISM IN PATIENTS UNDERGONG CHRONIC RENAL DIALYSIS

U-322 TREATMENT OF ALZHEIMER'S DEMENTIA

U-323 USE AS A BILE ACID SEQUESTRANT

U-324 METHOD OF TREATING AN ANIMAL, INCLUDING A HUMAN, SUFFERING FROM OR SUSCEPTIBLE TO PSYCHOSIS OR ACUTE MANIA EMPLOYING OLANZAPINE

U-325 METHOD OF TREATING A PATIENT SUFFERING FROM ANY OF A NUMBER OF LISTED CONDITIONS, INCLUDING "BIPOLAR DISORDER NOS" EMPLOYING OLANZAPINE

U-326 METHOD OF TREATING SCHIZOPHRENIA AND BIPOLAR DISORDER

U-327 METHOD OF TREATING A PATIENT SUFFERING FROM ANY OF A NUMBER OF LISTED PSYCHOTIC CONDITIONS EMPLOYING OLANZAPINE

U-328 METHOD OF TREATING A PATIENT SUFFERING FROM ANY OF A NUMBER OF LISTED CONDITIONS INCLUDING "A PSYCHOTIC CONDITION" EMPLOYING AN OLANZAPINE POLYMORPH

U-329 USE OF AVANDIA AS MONOTHERAPY, IN COMBINATION WITH METFORMIN, AND IN COMBINATION WITH SULFONYLUREAS TO IMPROVE GLYCEMIC CONTROL IN PATIENTS WITH TYPE 2 DIABETES MELLITUS

U-330 TREATMENT OF NAUSEA AND VOMITING

U-331 METHOD OF TREATING HYPERLIPIDEMIA WITH NICOTINIC ACID BY DOSING ONCE PER DAY IN THE EVENING OR AT NIGHT

U-332 TREATMENT OR PREVENTION OF BRONCHOSPASM

U-333 METHOD OF TREATING OCULAR HYPERTENSION

U-334 TREATMENT OF EXCESSIVE FEMALE FACIAL HAIR

U-335 USE OF PRAVASTATIN SODIUM FOR SECONDARY PREVENTION OF CORONARY EVENTS IN MEN AND WOMEN WHO HAVE HAD A MYOCARDIAL INFARCTION AND HAVE NORMAL CHOLESTEROL LEVELS

U-336 DIAGNOSTIC RADIOIMAGING

U-337 USE OF CARDIOLITE/MIRALUMA KIT FOR THE PREPARATION OF TC99M SESTAMIBI

U-338 METHODS FOR TREATING DISTURBANCES OF MOOD, DISTURBANCES OF APPETITE, DEPRESSED MOOD, OR CARBOHYDRATE CRAVING ALL ASSOCIATED WITH PREMENSTRUAL SYNDROME

U-339 PREVENTION OF CARDIO-TOXICITY CAUSED BY THE ADMINISTRATION OF DOXORUBICIN

U-340 THE LONG TERM TREATMENT OF GROWTH FAILURE DUE TO LACK OF ADEQUATE ENDOGENOUS GROWTH HORMONE SECRETION IN CHILDREN

U-341 METHOD FOR ENHANCING THE TREATMENT OF ... LATE LUTEAL PHASE DYSPHORIC DISORDER

U-342 METHOD FOR TREATMENT OF LATE LUTEAL PHASE DYSPHORIC DISORDER

U-343 REDUCTION OF INTESTINAL GAS, CRAMPING AND ANORECTAL IRRITATION

U-344 METHOD FOR INHIBITING HIV INFECTION BY ADMINISTERING RITONAVIR IN COMBINATION WITH ANOTHER HIV PROTEASE INHIBITOR

U-345 RITONAVIR AND ANOTHER HIV PROTEASE INHIBITOR FOR CONCOMITANT ADMINISTRATION FOR THE TREATMENT OF AN HIV INFECTION

U-346 METHOD FOR INHIBITING CYTOCHROME P450 MONOOXYGENASE WITH RITONAVIR AND A METHOD FOR IMPROVING THE PHARMACOKINETICS OF A DRUG THAT IS METABOLIZED BY CYTOCHROME P450 MONOOXYGENASE BY ADMIN THE DRUG AND RITONAVIR

U-347 METHOD OF USE IN COMBINATION WITH REVERSE TRANSCRIPTASE INHIBITORS

U-348 METHOD OF USE FOR INHIBITING HIV INFECTION

U-349 METHOD OF USE WHICH IS SUBJECT OF THE APPLICATION

U-350 PREPARATION OF A PHARMACEUTICAL COMPOSITION FOR CONCOMITANT ADMIN WITH A REVERSE TRANSCRIPTASE INHIBITOR

U-351 INHIBITING PROTEASE WITH LOPINAVIR AND INHIBITING AN HIV INFECTION WITH LOPINAVIR

U-352 INHIBITING HIV INFECTION BY ADMINISTERING RITONAVIR IN COMBINATION WITH A REVERSE TRANSCRIPTASE INHIBITOR

U-353 PREVENTION AND TREATMENT OF OSTEOPOROSIS

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

- U-354 METHOD OF TREATING HYPERLIPIDEMIA WITH NICOTINIC ACID WITHOUT CAUSING TREATMENT-LIMITING ELEVATIONS IN URIC ACID OR GLUCOSE LEVELS OR CAUSING LIVER DAMAGE, BY DOSING ONCE PER DAY IN THE EVENING OR AT NIGHT
- U-355 METHOD OF ASSISTING PERSON TO QUIT SMOKING...TRANSDERMALLY ADMIN NICOTINE VIA..PATCH ADHERED TO SKIN AT DOSING RATE APPROX SAME AS ABSORBED FROM SMOKING
- U-356 DELIVERING A MEDICINAL AEROSOL FORMULATION USING CFC-FREE PROPELLANT 134A.
- U-357 USE OF THE DRUG PRODUCT IN PHOTODYNAMIC THERAPEUTIC PROTOCOLS FOR THE TREATMENT OF AGE-RELATED MACULAR DEGENERATION AND RELATED CONDITIONS INVOLVING UNWANTED NEOVASCULATURE IN THE EYE
- U-358 DEPRESSION, OBSESSIVE COMPULSIVE DISORDER, PANIC DISORDER AND SOCIAL ANXIETY DISORDER
- U-359 METHOD OF USE OF VISICOL
- U-360 METHOD OF TREATING A PATIENT SUFFERING FROM ANY OF A NUMBER OF PATHOLOGICAL PSYCHOLOGICAL CONDITIONS INCLUDING MENTAL DISORDERS EMPLOYING OLANZAPINE AS PER THE INDICATION WHICH IS THE SUBJECT MATTER OF THIS SNDA-011
- U-361 MANAGEMENT OF ANXIETY DISORDERS AND THE SHORT-TERM RELIEF OF THE SYMPTOMS OF ANXIETY
- U-362 USE OF APPROVED FORMULATIONS TO TREAT ALL APPROVED DISEASE INDICATIONS
- U-363 METHOD OF TREATING A PATIENT SUFFERING FROM ANY OF A NUMBER OF PATHOLOGICAL PSYCHOLOGICAL CONDITIONS THAT RELATE TO THE USE OF A PSYCHOACTIVE SUBSTANCE EMPLOYING OLANZAPINE AS PER THE INDICATION THE SUBJECT MATTER OF SUPPLEMENT 011
- U-364 TREATING A PATIENT SUFFERING FROM OR SUSCEPTIBLE TO ANY NUMBER OF LISTED CONDITIONS INCLUDING PSYCHOSIS, EMPLOYING OLANZAPINE AS PER THE INDICATION WHICH IS THE SUBJECT MATTER OF THIS SNDA-011
- U-365 METHOD FOR THE TREATMENT OF CARDIOVASCULAR DISEASE THROUGH THE ADMINISTRATION OF A CALCIUM BLOCKING VASODILATOR IN OUR EXTENDED, CONTROLLED RELEASE FORMULATION
- U-366 METHOD FOR THE TREATMENT OF CARDIOVASCULAR DISEASE THROUGH THE ADMINISTRATION OF A CALCIUM BLOCKING VASODILATOR IN A DELAYED RELEASE FORMULATION
- U-367 TREATMENT OF CARDIOVASCULAR DISORDERS
- U-368 HEARTBURN
- U-369 METHOD OF CONTROLLING AND LOWERING INTRAOCULAR PRESSURE
- U-370 INTRAVAGINAL TREATMENT OF VAGINAL INFECTIONS WITH BUFFERED METRONIDAZOLE COMPOSITIONS
- U-371 APPROVAL FOR MARKETING ONLY UNDER A SPECIAL RESTRICTION PROGRAM APPROVED BY FDA CALLED "SYSTEM FOR THALIDOMIDE EDUCATION AND PRESCRIBING SAFETY" (S.T.E.P.S.)
- U-372 METHOD FOR ADMINISTERING A BENEFICIAL DRUG TO THE GI TRACT OF AN ANIMAL, WHICH METHOD COMPRISES ADMITTING AN OSMOTIC DEVICE ORALLY INTO THE ANIMAL...
- U-373 GENERAL USE CLAIM SUBMITTED FOR 12 NEXIUM PATIENTS STATING "PERTINENT TO THE CAPSULE FORMULATION FOR NEXIUM AND ITS INDICATIONS FOR THE TREATMENT OF GERD AND ERADICATION OF H.PYLORI TO REDUCE THE RISK OF DUODENAL ULCER RECURRENCE
- U-374 KIT ADAPTED AND DESIGNED TO PROVIDE BOTH DATA ON THE CURRENT REPRODUCTIVE STATUS OF A PATIENT AND CONTRACEPTION FOR THOSE WHO ARE NOT PREGNANT, BUT RECENTLY ENGAGED IN UNPROTECTED SEX
- U-375 METHOD OF USING RIBAVIRIN FOR TREATING A DISEASE RESPONSIVE TO RIBAVIRIN, E.G. HEPATITIS C
- U-376 TREATMENT OF INFLUENZA
- U-377 METHOD OF TREATING PT WITH CHRONIC HEPATITIS C HAVING HCV GENOTYPE 1 AND VIRAL LOAD GREATER THAN 2 MILLION COPIES/ML TO ERADICATE DETECTABLE HCV-RNA BY ADMIN COMBINATION OF RIBAVIRIN AND INTERFERON ALFA-2B FOR A LEAST 24 WEEKS
- U-378 METHOD FOR TREATING INCONTINENCE
- U-379 METHOD OF TREATING ONYCHOMYCOSIS
- U-380 COMBINATIONS OF TAXOL (PACLITAXEL) AND CISPLATIN WHICH ARE SUITABLE FOR THE TREATMENT OF OVARIAN AND NON-SMALL CELL LUNG CARCINOMAS
- U-381 TREATMENT OF HYPERPHOSPHATEMIA
- U-382 METHOD OF STABILIZING PROSTAGLANDIN
- U-383 METHOD FOR TREATING GLAUCOMA AND OCULAR HYPERTENSION

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-384 TREATMENT OF CMV RETINITIS

U-385 TREATMENT OF PEPTIC ULCERS

U-386 TREATMENT OF PATIENTS SUFFERING FROM A LATE ASTHMATIC REACTION OR LATE PHASE ASTHMA

U-387 TREATMENT OF PATIENTS WITH RESPIRATORY DISORDERS

U-388 SMOKING CESSATION AID APPLIED TO THE SKIN

U-389 SMOKING CESSATION AID APPLIED TO THE SKIN ON WAKING AND REMOVED PRIOR TO SLEEP AFTER ABOUT 16 HOURS

U-390 METHOD OF USING THE DRUG TO TREAT NEUROIMMUNOLOGIC DISEASES (INCLUDING MULTIPLE SCLEROSIS)

U-391 USE OF CASODEX IN COMBINATION WITH LHRH AGONISTS FOR THE TREATMENT OF PROSTATE CANCER

U-392 TREATMENT OF PATIENTS FOR INFLAMMATION

U-393 MANAGEMENT OF INCONTINENCE, MGT OF HORMONE REPLACEMENT THERAPY, TREATMENT OF INVOLUNTARY INCONTINENCE, MGT OVERACTIVE BLADDER AND INCREASING COMPLIANCE IN SUCH PT

U-394 METHOD OF USE OF ALPHAGAN

U-395 METHOD OF USE OF ALPHAGAN P

U-396 METHOD OF TREATING PEOPLE SUFFERING FROM DEPRESSION

U-397 METHOD OF TREATING PEOPLE SUFFERING FROM DEPRESSION WITHOUT AN INCREASE IN NAUSEA

U-398 TREATMENT OF GENERALIZED ANXIETY DISORDER

U-399 IN-THE-EYE USE OF CHLORINE DIOXIDE CONTAINING COMPOSITIONS

U-400 USE OF RIBAVIRIN TO INCREASE TYPE 1 CYTOKINE RESPONSE AND SUPPRESS TYPE 2 CYTOKINE RESPONSE TO LYMPHOCYTES, INCLUDING METHODS THAT TAKE ADVANTAGE OF SUCH MODULATION TO TREAT INFECTIONS AND INFESTATIONS

U-401 USE OF LOPINAVIR IN COMBINATION WITH REVERSE TRANSCRIPTASE INHIBITORS FOR TREATING HIV INFECTION AND IN COMBO WITH OTHER HIV PROTEASE INHIBITORS

U-402 TREATMENT OF ACTINIC KERATOSES

U-403 ANTI-ALLERGIC FOR VARIOUS ALLERGIC DISEASES

U-404 TREATMENT OF ALLERGIC CONJUNCTIVITIS

U-405 FOR WOMEN WITH SEVERE DIARRHEA-PREDOMINANT IRRITABLE BOWEL SYNDROME (IBS)

U-406 METHOD OF USE OF ATOVAQUONE AND PROGUANIL

U-407 METHOD OF TREATING OTOPATHY

U-408 FOR INDUCING OVULATION IN CONJUNCTION WITH A GONADOTROPIN RELEASING FACTOR ANTAGONIST AND RECRUITING OOCYTES FOR IN-VITRO FERTILIZATION

U-409 METHOD OF TREATING INFLAMMATION USING DRUG SUBSTANCE

U-410 METHOD OF REDUCING AMOUNT OF RESPECTIVE ACTIVE COMPONENTS ADMINISTERED TO A DIABETIC PATIENT BY ADMINISTERING A CHEMICAL COMPOUND HAVING A PARTICULAR FORMULA (INCLUDING PIOGLITAZONE) IN COMBINATION WITH AN INSULIN SECRETION ENHANCER

U-411 METHOD OF REDUCING THE SIDE EFFECTS OF ACTIVE COMPONENTS ADMINISTERED TO A DIABETIC PATIENT BY ADMINISTERING A CHEMICAL COMPOUND HAVING A PARTICULAR FORMULA (WHICH INCLUDES PIOGLITAZONE) IN COMBINATION WITH AN INSULIN PREPARATION

U-412 TREATMENT OF TYPE 2 DIABETES

U-413 USE OF THE ACTIVE INGREDIENT FOR INHIBITING THE BIOSYNTHESIS OF CHOLESTEROL AND TREATMENT OF ATHEROSCLEROSIS

U-414 A METHOD OF TREATING GLYCOMETABOLISM DISORDERS BY ADMINISTERING AN INSULIN SENSITIVITY ENHANCER (INCLUDING PIOGLITAZONE) IN COMBINATION WITH A BIGUANIDE

U-415 A METHOD FOR REDUCING THE AMOUNT OF ACTIVE COMPONENTS ADMINISTERED TO A DIABETIC PATIENT BY ADMINISTERING AN INSULIN SENSITIVITY ENHANCER (INCLUDING PIOGLITAZONE) IN COMBINATION WITH A BIGUANIDE AS SAID ACTIVE COMPONENTS

U-416 A METHOD FOR REDUCING SIDE EFFECTS OF ACTIVE COMPONENTS ADMINISTERED TO A DIABETIC PATIENT BY ADMINISTERING AN INSULIN SENSITIVITY ENHANCER (INCLUDING

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

- PIOGLITAZONE) IN COMBINATION WITH A BIGUANIDE AS SAID ACTIVE COMPONENTS
- U-417 COMBINATION USE OF AD-4833 WITH A BIGUANIDE
- U-418 A METHOD OF TREATING LIPID METABOLISM DISORDERS BY ADMINISTERING A CHEMICAL COMPOUND HAVING A PARTICULAR FORMULA (WHICH INCLUDES PIOGLITAZONE) IN COMBINATION WITH AN INSULIN SECRETION ENHANCER
- U-419 A METHOD OF TREATING LIPID METABOLISM DISORDERS BY ADMINISTERING AN INSULIN SENSITIVITY ENHANCER (INCLUDING PIOGLITAZONE) IN COMBINATION WITH A BIGUANIDE
- U-420 METHOD OF TREATMENT OF TYPE II DIABETES
- U-421 USE FOR SEDATION
- U-422 METHOD OF TREATING AT LEAST ONE OF ATTENTION DEFICIT DISORDER AND ATTENTION DEFICIT HYPERACTIVITY DISORDER
- U-423 METHOD OF TREATING AT LEAST ONE OF ATTENTION DEFICIT DISORDER, ATTENTION DEFICIT HYPERACTIVITY DISORDER, OR AIDS RELATED DEMENTIA
- U-424 FOR ONCE DAILY, BOLUS ADMINISTRATION TO A PATIENT IN ORDER TO ENGENDER TREATMENT FOR A NERVOUS DISORDER FOR SUBSTANTIALLY AN ENTIRE DAY ON A CHRONIC BASIS
- U-425 METHOD OF REDUCING SIDE EFFECTS OF ACTIVE COMPONENTS ADMIN TO A DIABETIC BY ADMIN A CHEMICAL COMPOUND HAVING FORMULA (INCL PIOGLITAZONE) IN COMBINATION WITH AN INSULIN SECRETION ENHANCER
- U-426 PREVENTION OF PREMATURE LH SURGES IN WOMEN UNDERGOING CONTROLLED OVARIAN STIMULATION
- U-427 METHOD OF TREATING ALLERGIC REACTIONS IN MAMMALS
- U-428 METHOD OF TREATING ALLERGY IN A MAMMAL USING THIS ACTIVE METABOLITE
- U-429 METHOD OF USING DESLORATADINE TO TREAT ALLERGIC RHINITIS
- U-430 METHOD OF TREATING A DIABETIC BY ADMINISTERING AN INSULIN SENSITIZER IN COMBINATION WITH AN INSULIN SECRETION ENHANCER, AND A DRUG PRODUCT COMPRISING AN INSULIN SENSITIZER AND AN INSULIN SECRETION ENHANCER
- U-431 POSTTRAUMATIC STRESS DISORDER
- U-432 REDUCTION OF ATHEROSCLEROTIC EVENTS (MYOCARDIAL INFARCTION, STROKE, AND VASCULAR DEATH) IN PATIENTS WITH ATHEROSCLEROSIS DOCUMENTED BY RECENT STROKE, RECENT MYOCARDIAL INFARCTION OR ESTABLISHED PERIPHERAL ARTERIAL DISEASE
- U-433 USE OF LEVOCARNITINE IN PREVENTION AND TREATMENT OF CARNITINE DEFICIENCY IN PATIENTS WITH END STAGE RENAL DISEASE WHO ARE UNDERGOING DIALYSIS
- U-434 CONTROLLED SYMPTOMS OF DIARRHEA, BLOATING PRESSURE AND CRAMPS, COMMONLY REFERRED TO AS GAS
- U-435 A TITRATION DOSING REGIMEN FOR THE TREATMENT OF PAIN USING AN INITIAL DOSE OF ABOUT 25MG
- U-436 ACUTE TREATMENT OF MIGRAINE ATTACKS WITH OR WITHOUT AURA IN ADULTS
- U-437 METHOD OF USE EQUAL TO PROCESS OF PREPARATION
- U-438 TREATMENT/PREVENTION OF NEURODEGENERATIVE DISEASE
- U-439 TREATMENT OF OBESITY
- U-440 METHOD FOR TRANSDERMAL ADMINISTRATION OF A DRUG THROUGH NON-SCROTAL SKIN USING A TRANSDERMAL DRUG DELIVERY DEVICE CONTAINING THE DRUG AND HAVING AN ADHESIVE SURFACE
- U-441 METHOD OF TREATING MS BY ADMINISTERING COPAXONE
- U-442 METHOD FOR DELIVERING A DRUG TO A PATIENT IN NEED OF THE DRUG, WHILE AVOIDING THE OCCURRENCE OF AN ADVERSE SIDE EFFECT KNOWN OR SUSPECTED OF BEING CAUSED BY SAID DRUG
- U-443 MANAGEMENT OF MODERATE TO SEVERE PAIN WHEN A CONTINUOUS, AROUND-THE-CLOCK ANALGESIC IS NEEDED FOR AN EXTENDED PERIOD OF TIME
- U-444 TREATMENT OF MIGRAINE
- U-445 USE AS AN ANTIMYCOTIC AGENT
- U-446 TOPICAL TREATMENT OF OCULAR HYPERTENSION AND GLAUCOMA
- U-447 METHOD OF TREATING HYPERLIPIDEMIA WITH NICOTINIC ACID BY DOSING ONCE PER DAY IN THE EVENING OR AT NIGHT

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

- U-448 METHOD OF TREATING HYPERLIPIDEMIA WITH NICOTINIC ACID WITHOUT CAUSING TREATMENT-LIMITING ELEVATIONS IN URIC ACID OR GLUCOSE LEVELS OR CAUSING LIVER DAMAGE, BY DOSING ONCE PER DAY IN THE EVENING OR AT NIGHT
- U-449 USE IN COMBINATION WITH 5-FLUOROURACIL AND LEUCOVORIN FOR THE TREATMENT OF METASTATIC COLORECTAL CANCER WHERE THE DOSE OF LEUCOVORIN IS AT LEAST 200MG PER SQUARE METER
- U-450 INTERMEDIATE REL NICOTINIC ACID FORMULATIONS HAVING UNIQUE URINARY METAB PROFILES RESULTING FROM ABSORPTION PROFILES OF NICOTINIC ACID FROM THE INTERMEDIATE NICOTINIC ACID FORMULATIONS, SUITABLE FOR TX HYPERLIPIDEMIA FOLLOWING QD DOSING
- U-451 TREATMENT OF DEPRESSION AND GENERALIZED ANXIETY DISORDER
- U-452 USE OF LANSOPRAZOLE FOR COMBATTING DISEASES CAUSED BY THE GENUS CAMPYLOBACTER (C.PYLORI=H.PYLORI)
- U-453 TREATMENT OF PLATELET ASSOCIATED ISCHEMIC DISORDERS
- U-454 METHOD OF TX A PT SUSPECTED OF HAVING HEPATITIS C BY ADMIN, IN COMBINATION, A CONJUGATE COMPRISING PEG 12000 & INTERFERON ALFA-2B IN AN AMT OF FROM 0.5MCG/KG TO 2MCG/KG, ONCE WEEKLY, AND RIBAVIRIN
- U-455 TREATMENT OF PULMONARY HYPERTENSION WITH UT-15
- U-456 METHOD OF DECREASING THE PRODUCTION OF A-BETA USING A COMPOSITION WHICH DECREASES BLOOD CHOLESTEROL IN PATIENTS AT RISK OF OR EXHIBITING SYMPTOMS OF ALZHEIMER'S DISEASE
- U-457 METHOD OF TREATING A VAGINAL FUNGAL INFECTION IN A FEMALE HUMAN
- U-458 METHOD OF USE OF IMAGENT
- U-459 TREATMENT OF DEPRESSION AND GENERALIZED ANXIETY DISORDER
- U-460 METHOD OF TREATING PSYCHIATRIC SYMPTOMS ASSOCIATED WITH PREMENSTRUAL DISORDERS USING SERTRALINE
- U-461 METHOD OF TREATMENT OF LATE LUTEAL PHASE DYSPHORIC DISORDER (PMDD) USING SERTRALINE
- U-462 SIGNS AND SYMPTOMS OF OSTEOARTHRITIS AND ADULT RHEUMATOID ARTHRITIS AND TREATMENT OF PRIMARY DYSMENORRHEA
- U-463 VENOGRAPHY
- U-464 PERIPHERAL ARTERIOGRAPHY
- U-465 CT IMAGING OF THE HEAD
- U-466 TREATMENT OF IRRITABLE BOWEL SYNDROME
- U-467 USE OF EPLERENONE IN COMBINATION WITH AN ANGIOTENSIN CONVERTING ENZYME (ACE) INHIBITOR FOR TREATING HYPERTENSION
- U-468 METHOD OF USING FEXOFENADINE HCL IN TREATING ALLERGIC RHINITIS
- U-469 TREATMENT OF GASTROESOPHAGEAL REFLEX DISEASE (GERD) AND ERADICATION OF H.PYLORI TO REDUCE RISK OF DUODENAL ULCER RECURRENCE
- U-470 THERAPY IN CHRONIC HEPATITIS B VIRUS INFECTION
- U-471 METHOD OF TREATING A PATIENT SUFFERING FROM DIABETES MELLITUS
- U-472 TREATMENT OF ATTENTION DEFICIT HYPERACTIVITY DISORDER USING METHYLPHENIDATE BI-MODAL RELEASE PROFILE EXTENDED-RELEASE CAPSULES
- U-473 TO REDUCE PLASMA CHOLESTEROL LEVELS IN A MAMMAL
- U-474 TO REDUCE PLASMA CHOLESTEROL LEVELS BY ADMIN EZETIMIBE IN COMBO WITH CHOLESTEROL BIOSYNTHESIS INHIB SELECTED FROM GROUP CONSISTING OF HMG COA REDUCTASE INHIBITORS INCL SIMVASTATIN
- U-475 TREATMENT OF CUTANEOUS MANIFESTATIONS OF CUTANEOUS T-CELL LYMPHOMA IN PATIENTS WHO ARE REFRACTORY TO AT LEAST ONE PRIOR SYSTEMIC THERAPY
- U-476 METHOD OF TREATING ANDROGEN RESPONSIVE/MEDIATED CONDITION IN MAMMAL BY ADMIN A SAFE, EFFECTIVE AMOUNT OF DUTASTERIDE OR PHARMACEUTICALLY ACCEPTABLE DERIVATIVE THEREOF..CONDITIONS INCLUDE BENIGN PROSTATIC HYPERTROPHY
- U-477 METHOD OF INHIBITING 5 ALPHA TESTOSTERONE REDUCTASE ENZYME WITH DUTASTERIDE OR ITS DERIVATIVE AND TREATING ANDROGEN RESPONSIVE/MEDIATED DISEASE INCLUDING BENIGN PROSTATIC HYPERPLASIA

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

- U-478 METHOD OF TREATING HEPATITIS C VIRAL INFECTION BY CONTINUOUS PARENTERAL ADMINISTRATION OF INTERFERON ALPHA 2-10 MILLION IU WEEKLY, SUBCUTANEOUSLY, INJECTION OF POLYMER-INTERFERON ALPHA CONJUGATE-POLYMER IS PEG-INTERFERON IS ALPHA 2B
- U-479 METHOD OF USING PEG-INTRON/REBETOL COMBINATION THERAPY AND INTRON/REBETOL COMBINATION THERAPY
- U-480 CONTRAST AGENT FOR MRI
- U-481 DISUBSTITUTED ACETYLENES BEARING HETEROAROMATIC AND HETEROBICYCLIC GROUPS HAVING RETINOID-LIKE ACTIVITY
- U-482 METHOD OF IN VITRO FERTILIZATION THERAPY INCLUDING MEANS FOR INDUCING OVULATION....
- U-483 METHOD FOR THE ADMINISTRATION OF DRUGS USING THAT COMPOUND
- U-484 METHOD OF TREATING A SKIN DISEASE WITH A CORTICOSTEROID-CONTAINING PHARMACEUTICAL COMPOSITION
- U-485 METHOD AND COMPOSITION FOR REDUCING NERVE INJURY PAIN ASSOCIATED WITH SHINGLES (HERPES ZOSTER AND POST-HERPETIC NEURALGIA)
- U-486 EXTERNAL PREPARATION FOR APPLICATION TO THE SKIN CONTAINING LIDOCAINE-DRUG RETAINING LAYER PLACED ON SUPPORT AND COMPRISES ADHESIVE GEL BASE 1-10% BY WEIGHT OF LIDOCAINE
- U-487 METHOD AND COMPOSITION FOR REDUCING NERVE INJURY PAIN ASSOCIATED WITH SHINGLES (HERPES ZOSTER AND POST-HERPETIC NEURALGIA)
- U-488 METHOD FOR REDUCING THE PAIN ASSOCIATED WITH HERPES-ZOSTER AND POST-HERPETIC NEURALGIA
- U-489 EXPECTORANT
- U-490 TESTOSTERONE REPLACEMENT THERAPY IN MALES FOR CONDITIONS ASSOCIATED WITH A DEFICIENCY OR ABSENCE OF ENDOGENOUS TESTOSTERONE
- U-491 METHOD OF DELIVERING A DRUG TO THE LUNG
- U-492 METHOD FOR THE TREATMENT OF SKIN, SUFFERING FROM A CONDITION SELECTED FROM A GROUP CONSISTING OF NONACNE INFLAMMATORY DERMATOSES... COMPRISING APPLYING TO AFFECTED AREA. A THERAPEUTICALLY EFFECTIVE AMT AZELAIC ACID
- U-493 TREATMENT OF TYPE 2 DIABETES MELLITUS
- U-494 TREATMENT OF ATTENTION-DEFICIT HYPERACTIVITY DISORDER
- U-495 PERITONEAL DIALYSIS SOLUTION
- U-496 METHOD FOR TREATING CHRONIC RENAL FAILURE
- U-497 RELIEF OF THE SIGNS AND SYMPTOMS OF OSTEOARTHRITIS AND RHEUMATOID ARTHRITIS
- U-498 INTRA-ARTERIAL AND INTRAVENOUS USES OF ULTRAVIST
- U-499 METHOD OF USING REBETOL CAPSULES IN COMBINATION WITH A CONJUGATE COMPRISING POLYETHYLENE GLYCOL (PEG) AND AN ALPHA INTERFERON, INCLUDING, FOR EXAMPLE, PEG-INTRON POWDER FOR INJECTION
- U-500 USE AS AN ANTIHYPERTENSIVE AGENT
- U-501 TREATMENT OF RECURRENT HERPES LABIALIS (COLD SORES) IN ADULTS
- U-502 PITYRIASIS VERSICOLOR
- U-503 GENERATOR MUST BE USED WITH INFUSION SYSTEM SPECIFICALLY LABELED FOR USE WITH GENERATOR
- U-504 TINEA PEDIS, TINEA CRURIS, TINEA CORPORIS
- U-505 ULTRASOUND CONTRAST AGENT
- U-506 PHARM PRODUCT CONTAINER 1ST CHAMBER IS DISPOSED AQUEOUS DILUENT SOL 2ND CHAMBER PHARM ACTIVE AGENT COMPRISING ACETYLCHOLINE, BUFFER IN 1ST CHAM IS SUFFICIENT TO BUFFER PH OF MIXED SOL RESULTING MIXTURE OF AQUEOUS DILUENT SOL & PHARM ACTIVE..
- U-507 ACROMEGALY IN PATIENTS W/INADEQUATE RESPONSE TO SURGERY AND/OR RADIATION THERAPY AND/OR MEDICAL THERAPIES, OR FOR WHOM THESE THERAPIES ARE NOT APPROPRIATE
- U-508 METHOD OF RELEASING 17-BETA OESTRADIOL PRECURSOR IN A SUBSTANTIALLY ZERO ORDER PATTERN FOR AT LEAST THREE WEEKS
- U-509 TREATMENT OF CUTANEOUS MANIFESTATIONS OF CUTANEOUS T-CELL LYMPHOMA IN PATIENTS WHO ARE REFRACTORY TO AT LEAST ONE PRIOR SYSTEMIC THERAPY

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-510 TOPICAL TREATMENT OF CUTANEOUS LESIONS IN PATIENTS WITH CUTANEOUS T-CELL LYMPHOMA (STAGE IA AND IB) WHO HAVE REFRACTORY OR PERSISTENT DISEASE AFTER OTHER THERAPIES OR WHO HAVE NOT TOLERATED OTHER THERAPIES

U-511 USE OF QUINOLONE COMPOUNDS AGAINST ANAEROBIC PATHOGENIC BACTERIA

U-512 USE OF QUINOLONE COMPOUNDS AGAINST ATYPICAL UPPER RESPIRATORY PATHOGENIC BACTERIA

U-513 METHODS OF USE OF ANTIMICROBIAL COMPOUNDS AGAINST PATHOGENIC MYCOPLASMA BACTERIA

U-514 PREVENTION OF OVULATION IN A WOMAN

U-515 TREATMENT OF MULTIPLE MYELOMA PATIENTS WHO HAVE RECEIVED AT LEAST TWO PRIOR THERAPIES AND HAVE DEMONSTRATED DISEASE PROGRESSION ON THE LAST THERAPY

U-516 METHOD OF TREATING A PSYCHOTIC DISEASE

U-517 STABLE GEL FORMULATION FOR TOPICAL TREATMENT OF SKIN CONDITIONS

U-518 OBSESSIVE COMPULSIVE DISORDER

U-519 POST OPERATIVE NAUSEA AND VOMITING

U-520 PREMENOPAUSAL OSTEOPOROSIS

U-521 METHOD OF USING RIBAVIRIN IN COMBINATION WITH INTRON A (INTERFERON ALPHA-2 B RECOMBINANT) INJECTION TO TREAT PATIENTS WITH CHRONIC HEPATITIS C

U-522 TREATMENT OF CMV RETINITIS BY INTRAVITREAL ADMIN OF A PHOSPHOROTHIOATE OLIGONUCLEOTIDE CAPABLE OF HYBRIDIZING WITH CMV MRNA

U-523 METHOD OF TREATING INFECTION BY CRYPTOSPORIDIUM PARVUM IN AN IMMUNOCOMPROMISED MAMMAL

U-524 METHOD OF TREATING DIARRHEA

U-525 METHOD OF TREATING PARASITIC INFECTIONS

U-526 METHOD OF PROVIDING CONTROLLED RELEASE OF A TREATING AGENT USING A CONTROLLED RELEASE COMPOSITION

U-527 METHOD OF DELIVERING AN ACTIVE INGREDIENT USING A PROGRESSIVE HYDRATION BIOADHESIVE

U-528 PREVENTION OF CHEMOTHERAPY-INDUCED NAUSEA AND VOMITING

U-529 ONCE DAILY TREATMENT OF ASTHMA WITH NEBULIZED BUDESONIDE

U-530 TREATMENT OF HERPES ZOSTER, TREATMENT OF GENITAL HERPES, TREATMENT OF COLD SORES, SUPPRESSION OF GENITAL HERPES IN IMMUNOCOMPETENT AND HIV-INFECTED INDIVIDUALS, REDUCTION OF RISK OF HETEROSEXUAL TRANSMISSION OF GENITAL HERPES

U-531 TREATMENT OF PATIENTS WITH ESSENTIAL HYPERTENSION. MAY BE USED ALONE OR GIVEN WITH OTHER CLASSES OF ANTIHYPERTENSIVES, ESPECIALLY THIAZIDE DERIVATIVES

U-532 TREATMENT OF BRONCHOSPASM ASSOCIATED WITH COPD IN PATIENTS REQUIRING MORE THAN ONE BRONCHO DILATOR

U-533 ERECTILE DYSFUNCTION

U-534 HUMALOG IS AN INSULIN ANALOG THAT IS INDICATED IN THE TREATMENT OF PATIENTS WITH DIABETES MELLITUS FOR THE CONTROL OF HYPERGLYCEMIA

U-535 TREATMENT OF SOCIAL ANXIETY DISORDER

U-536 CONTRAST AGENT FOR MAGNETIC RESONANCE IMAGING

U-537 TREATMENT OF CONDITIONS RELATED TO HYPERALDOSTERONISM SUCH AS HYPERTENSION AND CARDIAC INSUFFICIENCY, WITH EPLERENONE

U-538 FIRST LINE TREATMENT OF SEVERE HYPERTENSION, IN PATIENTS WITH HYPERTENSION SEVERE ENOUGH THAT THE VALUE OF ACHIEVING PROMPT BLOOD PRESSURE CONTROL EXCEEDS THE RISK OF INITIATING COMBINATION THERAPY IN THESE PATIENTS

U-539 TREATMENT OF MODERATE TO SEVERE DEMENTIA OF THE ALZHEIMER'S TYPE

U-540 TREATMENT OF FUNGAL INFECTIONS

U-541 METHOD OF TREATMENT OF ADULTS INFECTED WITH HIV-1

U-542 METHOD OF TREATING PATIENT WITH TYPE 2 DIABETES BY ONCE DAILY ADMINISTRATION

U-543 TREATMENT OF SCHIZOPHRENIA

U-544 TREATMENT OF OVERACTIVE BLADDER. TREATMENT OF URINARY INCONTINENCE.

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

- U-545 METHOD FOR THE PREVENTION AND/OR TREATMENT OF THROMBOTIC EPISODES, SUCH AS MYOCARDIAL INFARCTION, IN A HUMAN PATIENT AND METHOD FOR THE PREVENTION OF VENOUS THROMBOSIS IN A POSTOPERATIVE HUMAN PATIENT
- U-546 USE OF REPAGLINIDE IN COMBINATION WITH METFORMIN TO LOWER BLOOD GLUCOSE
- U-547 MAINTENANCE MONOTHERAPY FOR BIPOLAR DISORDER
- U-548 A METHOD OF REDUCING FLUSH IN AN INDIVIDUAL BEING TREATED FOR A LIPIDEMIC DISORDER AND EFFECTIVELY TREATING THE LIPIDEMIC DISORDER
- U-549 USE IN THE TREATMENT OF MEN WITH ADVANCED SYMPTOMATIC PROSTATE CANCER
- U-550 TREATMENT OF BIPOLAR DISORDER AND SCHIZOPHRENIA
- U-551 METHOD FOR REDUCING TOXICITY OF ALIMTA TREATED PATIENTS BY ADMINISTERING FOLIC ACID
- U-552 TREATMENT OF HYPERTENSION AND HYPERLIPIDEMIA WITH A SINGLE COMPOSITION
- U-553 MANAGEMENT OF PAIN AND DISCOMFORT ASSOCIATED WITH PERIDONTAL SCALING AND ROOT PLANNING PROCEDURES BY APPLICATION OF AN EUTECTIC MIXTURE OF LOCAL ANESTHETICS TO PERIDONTAL POCKETS
- U-554 TREATING HIV INFECTION WITH INDINAVIR SULFATE IN COMBINATION WITH ANTIRETROVIRAL AGENTS
- U-555 TREATMENT OF COMPLICATED URINARY TRACT INFECTIONS AND ACUTE UNCOMPLICATED PYELONEPHRITIS
- U-556 USE AS ADJUNCT DIAGNOSTIC FOR SERUM THYROGLOBULIN (TG) TESTING
- U-557 NASAL TREATMENT OF SEASONAL AND PERENNIAL ALLERGIC RHINITIS SYMPTOMS
- U-558 INDICATED FOR THE RELIEF OF BRONCHOSPASM IN PATIENTS 2-12 YEARS OF AGE WITH ASTHMA (REVERSIBLE OBSTRUCTIVE AIRWAY DISEASE)
- U-559 METHOD OF DECREASING OR REDUCING PARATHYROID HORMONE LEVEL; METHOD OF MODULATING PARATHYROID HORMONE SECRETION;METHOD OF TREATING HYPERPARATHYROIDISM; METHOD OF REDUCING SERUM IONIZED CALCIUM LEVEL
- U-560 METHOD OF DECREASING PARATHYROID HORMONE LEVEL;METHOD OF TREATING HYPERPARATHYROIDISM
- U-561 COSOPT IS INDICATED FOR THE REDUCTION OF ELEVATED INTRAOCULAR PRESSURE IN PATIENTS WITH OPEN-ANGLE GLAUCOMA OR OCULAR HYPERTENSION WHO ARE INSUFFICIENTLY RESPONSIVE TO BETA BLOCKERS
- U-562 TOPICAL TREATMENT OF CUTANEOUS LESIONS IN PATIENTS WITH AIDS-RELATED KAPOSI'S SARCOMA
- U-563 MARINOL IS INDICATED FOR, INTER ALIA, ANOREXIA ASSOCIATED WITH WEIGHT LOSS IN PATIENTS WITH AIDS
- U-564 TREATMENT OF HIV IN CONCOMITANT THERAPY
- U-565 TREATMENT OF SEASONAL AND PERENNIAL ALLERGIC RHINITIS SYMPTOMS, AND CHRONIC URTICARIA
- U-566 FOR THE LONG-TERM, ONCE-DAILY, MAINTENANCE TREATMENT OF BRONCHOSPASM ASSOCIATED WITH CHRONIC OBSTRUCTIVE PULMONARY DISEASE (COPD), INCLUDING CHRONIC BRONCHITIS AND EMPHYSEMA
- U-567 METHOD OF TREATING INFERTILITY
- U-568 METHOD OF USING FSH ALONE (WITHOUT EXOGENOUS LH) IN IN VITRO FERTILIZATION
- U-569 METHOD OF USING FSH ALONE (WITHOUT EXOGENOUS LH) IN IN VITRO FERTILIZATION AND WHEREIN THEREAFTER AN OVULATORY INDUCING AMOUNT OF HCG IS ADMINISTERED
- U-570 METHOD OF USING FSH ALONE (WITHOUT EXOGENOUS LH) IN IN VITRO FERTILIZATION AND WHEREIN THE DAILY AMOUNT OF FSH IS ABOUT 5-10 IU/KG
- U-571 TREATMENT OF AGITATION ASSOCIATED WITH SCHIZOPHRENIA AND BIPOLAR I MANIA
- U-572 INTENSIVE CARE UNIT SEDATION
- U-573 TREATMENT OF ACUTE PROMYELOGENOUS LEUKEMIA (APL)
- U-574 PROPHYLAXIS AND TREATMENT OF THE NASAL SYMPTOMS OF SEASONAL ALLERGIC RHINITIS AND TREATMENT OF THE NASAL SYMPTOMS OF PERENNIAL ALLERGIC RHINITIS IN ADULTS AND PEDIATRIC PATIENTS 12 YEARS OF AGE AND OLDER
- U-575 LOTEMAX OPHTHALMIC SUSPENSION IS INDICATED FOR THE TREATMENT OF STEROID RESPONSIVE CONDITIONS OF THE PALPEBRAL BULBAR CONJUNCTIVA, CORNEA AND ANTERIOR

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

SEGMENT OF THE GLOBE.

- U-576 ALREX OPHTHALMIC SUSPENSION IS INDICATED FOR THE TEMPORARY RELIEF OF THE SIGNS AND SYMPTOMS OF SEASONAL ALLERGIC CONJUNCTIVITIS.
- U-577 TREATMENT OF BENIGN PROSTATIC HYPERPLASIA WITH FINASTERIDE IN COMBINATION WITH DOXAZOSIN
- U-578 TREATMENT OF COMMUNITY ACQUIRED PNEUMONIA, ACUTE EXACERBATION OF CHRONIC BRONCHITIS, AND ACUTE BACTERIAL SINUSITIS CAUSED BY SUSCEPTIBLE STRAINS OF DESIGNATED MICROORGANISMS IN PATIENTS 18 YEARS AND OLDER.
- U-579 TREATMENT OF EPILEPSY AND/OR MIGRAINE.
- U-580 TREATMENT OF DISORDERS OF THE SEROTONERGIC SYSTEM SUCH AS DEPRESSION AND ANXIETY-RELATED DISORDERS
- U-581 METHOD OF TREATING A CONDITION CAPABLE OF TREATMENT BY INHALATION, E.G. ASTHMA, COMPRISING ADMINISTRATION OF A FORMULATION CLAIMED IN US PATENT NO. 6743413
- U-582 METHOD FOR THE TREATMENT OF A RESPIRATORY DISORDER, E.G. ASTHMA, COMPRISING ADMINISTERING AN EFFECTIVE AMOUNT OF AN AEROSOL COMPOSITION TO A PATIENT FROM A METERED DOSE INHALER SYSTEM AS CLAIMED IN US PATENT NO. 6253762
- U-583 METHOD FOR THE TREATMENT OF A RESPIRATORY DISORDER, E.G. ASTHMA, COMPRISING ADMINISTERING TO A PATIENT BY INHALATION, A METERED AEROSOL DOSE OF A DRUG FORMULATION FROM THE METERED DOSE INHALER SYSTEM CLAIMED IN US 6546928
- U-584 SINGLE-DOSE ADMINISTRATION BY THE EPIDURAL ROUTE, AT THE LUMBAR LEVEL, FOR THE TREATMENT OF PAIN FOLLOWING MAJOR SURGERY
- U-585 TO PROMOTE WEIGHT GAIN AFTER WEIGHT LOSS IN CERTAIN TYPES OF PATIENTS
- U-586 AN INTERMEDIATE RELEASE NICOTINIC ACID FORMULATION SUITABLE FOR ORAL ADMINISTRATION ONCE-A-DAY AS A SINGLE DOSE FOR TREATING HYPERLIPIDEMIA WITHOUT CAUSING DRUG-INDUCED HEPATOTOXICITY OR ELEVATIONS IN URIC ACID OR GLUCOSE OR BOTH
- U-587 USE OF EPLERENONE IN COMBINATION WITH AN ANGIOTENSIN CONVERTING ENZYME (ACE) INHIBITOR (AND OPTIONALLY A DIURETIC) FOR TREATING CONGESTIVE HEART FAILURE AND HYPERTENSION
- U-588 SHORT-TERM TREATMENT OF ACTIVE DUODENAL ULCER; TREATMENT OF HEARTBURN AND OTHER SYMPTOMS ASSOCIATED WITH GERD; SHORT-TERM TREATMENT OF EROSIIVE ESOPHAGITIS; MAINTENANCE OF HEALING OF EROSIIVE ESOPHAGITIS
- U-589 METHOD FOR TREATMENT OF A RESPIRATORY DISORDER, E.G., BRONCHOSPASM, COMPRISING ADMINISTERING AN EFFECTIVE AMOUNT OF AN AEROSOL COMPOSITION TO A PATIENT FROM A METERED DOSE INHALER SYSTEM AS CLAIMED IN U.S. PATENT NO. 6131966
- U-590 METHOD FOR TREATMENT OF A RESPIRATORY DISORDER, E.G., BRONCHOSPASM, COMPRISING ADMINISTERING TO A PATIENT BY ORAL OR NASAL INHALATION A DRUG FORMULATION BY USING THE METERED DOSE INHALER SYSTEM AS CLAIMED IN US PATENT NO. 6532955
- U-591 TREATMENT OF ATTENTION DEFICIT HYPERACTIVITY DISORDER USING A DOSAGE FORM WHICH PROVIDES ONCE-DAILY ORAL ADMINISTRATION OF A PHENIDATE DRUG
- U-592 TREATMENT OF PRIMARY HYPERCHOLESTEROLEMIA, MIXED HYPERLIPIDEMIA AND/OR HOMOZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA (HOFH)
- U-593 TREATMENT OF PRIMARY HYPERCHOLESTEROLEMIA, MIXED HYPERLIPIDEMIA AND/OR HOMOZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA (HOFH)
- U-594 PREVENTION OF POSTMENOPAUSAL OSTEOPOROSIS
- U-595 35 MG ORALLY ONCE A WEEK FOR PREVENTION OF OSTEOPOROSIS IN POSTMENOPAUSAL WOMEN; 35 MG ORALLY ONCE A WEEK FOR TREATMENT OF OSTEOPOROSIS IN POSTMENOPAUSAL WOMEN
- U-596 TREATMENT OF HORMONE RECEPTOR POSITIVE METASTATIC BREAST CANCER IN POSTMENOPAUSAL WOMEN WITH DISEASE PROGRESSION FOLLOWING ANTIESTROGEN THERAPY
- U-597 FORTEO IS INDICATED FOR THE TREATMENT OF POST MENOPAUSAL WOMEN WITH OSTEOPOROSIS WHO ARE AT HIGH RISK FOR FRACTURE
- U-598 PROPHYLACTIC TREATMENT OF MIGRAINE
- U-599 METHOD FOR TREATING ALLERGIC CONJUNCTIVITIS
- U-600 A METHOD OF TREATING A PATIENT IN NEED OF OPHTHALMIC ANTIMICROBIAL THERAPY WITH LEVOFLOXACIN
- U-601 TREATMENT OF BIPOLAR DISORDER
- U-602 SIGNS AND SYMPTOMS OF OSTEOARTHRITIS, RHEUMATOID ARTHRITIS IN ADULTS, AND/OR

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

- PAUCIARTICULAR OR POLYARTICULAR COURSE JUVENILE RHEUMATOID ARTHRITIS, ACUTE PAIN IN ADULTS; PRIMARY DYSMENORRHEA; AND/OR ACUTE MIGRAINE ATTACKS IN ADULTS
- U-603 METHOD OF TREATING INFECTIONS COMPRISING ORALLY ADMINISTERING AN EFFECTIVE AMOUNT OF THE FDA APPROVED ORAL SUSPENSION
- U-604 METHOD OF LOWERING BLOOD GLUCOSE BY ONCE DAILY ADMINISTRATION
- U-605 TREATMENT OF MAJOR DEPRESSIVE DISORDER (MDD); ALTHOUGH THE MECHANISM OF THE ANTIDEPRESSANT ACTION OF DULOXETINE IN HUMANS IS UNKNOWN, IT IS BELIEVED TO BE RELATED TO ITS POTENTIATION OF SERATONERGIC AND NORADRENERGIC ACTIVITY IN THE CNS
- U-606 USE OF IRINOTECAN IN COMBINATION WITH 5-FLUOROURACIL AND LEUCOVORIN FOR THE TREATMENT OF METASTATIC COLORECTAL CANCER
- U-607 CANCIDAS IS INDICATED FOR EMPIRICAL THERAPY FOR PRESUMED FUNGAL INFECTIONS IN FEBRILE, NEUTROPENIC PATIENTS.
- U-608 USE OF QUINOLONE COMPOUNDS AGAINST PNEUMOCOCCAL PATHOGENIC BACTERIA
- U-609 USE OF QUINOLONE COMPOUNDS AGAINST QUINOLONE-RESISTANT PNEUMOCOCCAL PATHOGENIC BACTERIA
- U-610 ATROVENT HFA (IPRATROPIUM BROMIDE HFA) INHALATION AEROSOL IS INDICATED AS A BRONCHODILATOR FOR MAINTENANCE TREATMENT OF BRONCHOSPASM ASSOCIATED WITH CHRONIC OBSTRUCTIVE PULMONARY DISEASE, INCLUDING CHRONIC BRONCHITIS AND EMPHYSEMA.
- U-611 METHOD OF USING DESLORATADINE TO TREAT SEASONAL AND PERENNIAL ALLERGIC RHINITIS, PRURITIS, AND CHRONIC IDIOPATHIC URTICARIA IN PATIENTS 2 YEARS OF AGE AND OLDER
- U-612 TREATMENT OF SEASONAL ALLERGY SYMPTOMS WITH NASAL CONGESTION IN ADULTS AND CHILDREN 12 YEARS OF AGE AND OLDER
- U-613 REDUCTION OF SERUM PHOSPHATE
- U-614 TREATMENT OF SEXUAL DYSFUNCTION
- U-615 ADJUNCTIVE THERAPY TO DIET IN ADULTS TO REDUCE LDL-C, TOTAL-C, TRIGLYCERIDES AND APO B, AND INCREASE HDL-C IN PATIENTS WITH PRIMARY HYPERCHOLESTEROLEMIA OR MIXED DYSLIPIDEMIA (TYPES IIA, IIB) AND TO TREAT HYPERTRIGLYCERIDEMIA (TYPES IV, V)
- U-616 MANAGEMENT OF PERSISTENT, MODERATE TO SEVERE PAIN IN PATIENTS REQUIRING CONTINUOUS, AROUND-THE-CLOCK ANALGESIA WITH A HIGH POTENCY OPIOID FOR AN EXTENDED PERIOD OF TIME GENERALLY WEEKS TO MONTHS OR LONGER
- U-617 TREATMENT OF ACUTE PROMYELOGENOUS LEUKEMIA (APL)
- U-618 USE OF ROSUVASTATIN CALCIUM TO REDUCE ELEVATED TOTAL-C, LDL-C, APOB, NONHDL-C OR TG LEVELS; TO INCREASE HDL-C IN ADULT PATIENTS WITH PRIMARY HYPERLIPIDEMIA OR MIXED DYSLIPIDEMIA; AND TO SLOW THE PROGRESSION OF ATHEROSCLEROSIS.
- U-619 TREATMENT OF MALIGNANT NEOPLASM
- U-620 TREATMENT OF INSOMNIA
- U-621 METHOD OF TREATING CANCER
- U-622 TREATMENT OF VEGF MEDIATED OCULAR DISEASE.
- U-623 SHORT TERM TREATMENT OF ACTIVE BENIGN GASTRIC ULCER
- U-624 REDUCTION OF RISK OF UPPER GASTROINTESTINAL BLEEDING IN CRITICALLY ILL PATIENTS
- U-625 ALLERGIC RHINITIS OR NASAL POLYPS
- U-626 CLOLAR IS INDICATED FOR THE TREATMENT OF PEDIATRIC PATIENTS 1 TO 21 YEARS OLD WITH RELAPSED OR REFRACTORY ACUTE LYMPHOBLASTIC LEUKEMIA AFTER AT LEAST TWO PRIOR REGIMENS
- U-627 TREATMENT OF PATIENTS USING EXTENDED-RELEASE CARBAMAZEPINE
- U-628 USE OF AVANDIA IN COMBINATION WITH A SULFONYLUREA, AND IN COMBINATION WITH METFORMIN AND A SULFONYLUREA TO IMPROVE GLYCEMIC CONTROL IN PATIENTS WITH TYPE 2 DIABETES MELLITUS
- U-629 METHOD OF INDUCING A HYPNOTIC OR SEDATIVE EFFECT IN A HUMAN BY ADMINISTERING ESZOPICLONE
- U-630 TREATING URINARY INCONTINENCE BY ADMINISTERING AN EXTENDED-RELEASE FORM OF DARIFENACIN
- U-631 TREATING A DISEASE OF ALTERED MOTILITY OR TONE OF SMOOTH MUSCLE BY ADMINISTERING A MUSCARINIC RECEPTOR ANTAGONIZING AMOUNT OF DARIFENACIN

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-632 METHOD OF TREATMENT OF CANCER BY ADMINISTERING PARTICLES OF PACLITAXEL THAT HAVE A PROTEIN COATING

U-633 METHOD FOR TREATMENT OF TUMORS BY ADMINISTERING PACLITAXEL AT A DOSE IN THE RANGE OF ABOUT 30MG/METER SQUARE TO ABOUT 100MG/METER SQUARE IN A PHARMACEUTICALLY ACCEPTABLE FORMULATION THAT DOES NOT CONTAIN CREMOPHOR

U-634 METHOD FOR DELIVERY OF A BIOLOGIC (INCLUDING ANTINEOPLASTIC AGENTS) BY ADMINISTERING TO A PATIENT AN EFFECTIVE AMOUNT OF A BIOLOGIC AS A SOLID OR LIQUID WITH A POLYMERIC BIOCOMPATIBLE MATERIAL

U-635 TREATMENT OF GERD, MAINTENANCE OF HEALING OF EROSIIVE ESOPHAGITIS AND RISK REDUCTION OF NSAID ASSOCIATED GASTRIC ULCERS

U-636 TREATMENT OR PREVENTION OF BRONCHOSPASM OR ASTHMATIC SYMPTOMS

U-637 TREATMENT OF DIABETES WITH AN AMYLIN AGONIST

U-638 TREATMENT OF DIABETES WITH AN AMYLIN AGONIST, INCLUDING WITH INSULIN

U-639 TREATMENT OF A MAMMAL HAVING A NEED OF OR REDUCED ABILITY TO PRODUCE INSULIN WITH AN INSULIN AND AN AMYLIN SUCH AS PRAMLINTIDE

U-640 USE OF AN AMYLIN AGONIST TO REDUCE GASTRIC MOTILITY AND TREAT POST PRANDIAL HYPERGLYCEMIA

U-641 USE OF AN AMYLIN AGONIST HAVING SPECIFIED BINDING ACTIVITY TO REDUCE GASTRIC MOTILITY, INCLUDING USE THROUGH PARENTERAL ADMINISTRATION

U-642 TREATMENT AND PREVENTION OF OSTEOPOROSIS

U-643 THE SHORT TERM TREATMENT (UP TO 10 DAYS) IN PTS HAVING GASTROESOPHAGEAL REFLUX DISEASE (GERD) AS AN ALTERNATIVE TO ORAL THERAPY IN PTS WHEN THERAPY WITH NEXIUM CAPSULES IS NOT POSSIBLE OR APPROPRIATE

U-644 TREATMENT OF SEASONAL ALLERGIC RHINITIS

U-645 TREATMENT OF ASTHMA

U-646 METHOD OF TREATING OTITIS

U-647 TREATMENT OF OSTEOPOROSIS IN POST MENOPAUSAL WOMEN AND/OR THE TREATMENT TO INCREASE BONE MASS IN MEN WITH OSTEOPOROSIS

U-648 THE TREATMENT OF OSTEOPOROSIS IN POSTMENOPAUSAL WOMEN AND/OR THE TREATMENT TO INCREASE BONE MASS IN MEN

U-649 A METHOD FOR TREATING A TUMOR DISEASE

U-650 TREATMENT OF ESOPHAGEAL CANDIDIASIS AND PROPHYLAXIS OF CANDIDA INFECTIONS IN HSCT PATIENTS

U-651 TREATMENT OF ACUTE PROMYELOCYTIC LEUKEMIA (APL)

U-652 TREATMENT OF CARDIAC ARRHYTHMIA

U-653 STIMULATING INSULIN RELEASE BY ADMINISTERING EXENATIDE

U-654 LOWERING PLASMA GLUCAGON IN A SUBJECT IN NEED THEREOF, INCLUDING ONE WITH TYPE 2 DIABETES, BY ADMINISTERING AN EXENDIN OR ANALOG, SUCH AS EXENDIN-4

U-655 TREATMENT OF MILD TO MODERATE ACTIVE CHROHN'S DISEASE INVOLVING THE ILEUM AND/OR THE ASCENDING COLON AND THE MAINTENANCE OF CLINICAL REMISSION OF MILD TO MODERATE CROHN'S DISEASE INVOLVING THE ILEUM AND/OR ASCENDING COLON FOR UP TO 3 MONTHS

U-656 REDUCING GASTRIC MOTILITY OR DELAYING GASTRIC EMPTYING BY ADMINISTERING AN EXENDIN, SUCH AS EXENDIN-4

U-657 PREVENTION OF OSTEOPOROSIS IN POSTMENOPAUSAL WOMEN

U-658 TREATMENT OF ADVANCED HORMONE-DEPENDENT BREAST CANCER

U-659 TREATMENT OF LOCALLY ADVANCED OR METASTATIC NON SMALL-CELL LUNG CANCER (NSCLC) AFTER FAILURE OF AT LEAST ONE PRIOR CHEMOTHERAPY REGIMEN

U-660 TREATMENT OF HYPERTENSION AND TREATMENT OF HEART FAILURE

U-661 TREATMENT OF SEIZURE DISORDER

U-662 TREATMENT OF OSTEOPOROSIS IN POSTMENOPAUSAL WOMEN

U-663 THE TREATMENT OF UNCOMPLICATED URINARY TRACT INFECTIONS

U-664 TREATMENT OF CONDITIONS FOR WHICH AN ALDOSTERONE RECEPTOR BLOCKER IS INDICATED, SUCH AS HYPERTENSION, HEART FAILURE, AND POST-MYOCARDIAL INFARCTION

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-665 METHOD OF USING THE DRUG SUBSTANCE/DRUG PRODUCT FOR ULTRASOUND IMAGING

U-666 METHOD OF TREATING ADHD

U-667 MANAGEMENT OF INCONTINENCE; METHOD FOR TREATING INCONTINENCE

U-668 LEVEMIR IS A LONG-ACTING BASAL INSULIN ANALOG THAT IS INDICATED IN THE TREATMENT OF PATIENTS WITH DIABETES MELLITUS

U-669 INDICATION OF TYPE II DIABETES

U-670 TREATMENT OF HIV-1 INFECTION BY THE CO-ADMINISTRATION OF TIPRANAVIR AND RITONAVIR.

U-671 PREVENTION AND TREATMENT OF SECONDARY HYPERPARATHYROIDISM ASSOCIATED WITH CHRONIC KIDNEY DISEASE (CKD) STAGE 3 AND 4

U-672 TREATMENT OF INFLAMMATION OR AN INFLAMMATION-ASSOCIATED DISORDER

U-673 METHOD OF TREATMENT WITH ONCE-DAILY DOSES OF 625MG/5ML

U-674 METHOD OF TREATING INSOMNIA CHARACTERIZED BY DIFFICULTY WITH SLEEP ONSET

U-675 PROPHYLAXIS AND CHRONIC TREATMENT OF ASTHMA; RELIEF OF SYMPTOMS OF ALLERGIC RHINITIS

U-676 METHOD OF TREATING ATTENTION DEFICIT DISORDER USING ORAL ADMINISTRATION OF A BI-MODAL OR PULSATILE RELEASE COMPOSITION

U-677 A METHOD OF TREATING DISEASE AMENABLE TO TREATMENT WITH A PHENIDATE DRUG BY ONCE DAILY ORAL ADMINISTRATION OF AN EXTENDED RELEASE DOSAGE FORM

U-678 METHOD OF TREATING ATTENTION DEFICIT DISORDER AND/OR ATTENTION DEFICIT HYPERACTIVITY DISORDER

U-679 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN PATIENTS WITH TYPE 2 DIABETES WHO ARE ALREADY TREATED WITH A PIOGLITAZONE AND METFORMIN

U-680 A METHOD OF TREATING DYSLIPIDEMIA AND DYSLIPOPROTEINEMIA USING A DOSAGE FORM THAT CAN PROVIDE AN EFFECTIVE AMOUNT OF FENOFIBRATE TO A PATIENT IN A FASTED STATE WHICH IS AT LEAST 90% OF THE AUC AMOUNT PROVIDED BY THE DOSAGE FORM

U-681 TREATMENT OF PRIMARY IGF-1 DEFICIENCY

U-682 NON-BENZODIAZEPINE HYPNOTIC AGENT INDICATED FOR TREATMENT OF INSOMNIA, CHARACTERIZED BY DIFFICULTIES WITH SLEEP ONSET AND/OR SLEEP MAINTENANCE

U-683 PREVENTION OR TREATMENT OF ISCHEMIC HEART DISEASE

U-684 TREATMENT OF UNCOMPLICATED SKIN MANIFESTATIONS OF CHRONIC IDIOPATHIC URTICARIA IN ADULTS AND CHILDREN 6 YEARS OF AGE AND OLDER

U-685 EXPECTORANT AND COUGH SUPPRESSANT

U-686 EXPECTORANT AND NASAL DECONGESTANT

U-687 REDUCING FOOD INTAKE IN A SUBJECT WITH TYPE 2 DIABETES BY ADMINISTERING AN EXENDIN, SUCH AS EXENDIN-4

U-688 TREATMENT OF HIV-INFECTION IN COMBINATION WITH OTHER ANTIRETROVIRAL AGENTS

U-689 TREATMENT OF PATIENTS WITH T-CELL ACUTE LYMPHOBLASTIC LEUKEMIA WHOSE DISEASE HAS NOT RESPONDED TO OR HAS RELAPSED FOLLOWING TREATMENT WITH AT LEAST TWO CHEMOTHERAPY REGIMENS

U-690 TO IMPROVE GLYCEMIC CONTROL IN PATIENTS WITH TYPE 2 DIABETES MELLITUS

U-691 USE AS A MONOTHERAPY, IN COMBINATION WITH A SULFONYLUREA, METFORMIN OR INSULIN OR IN COMBINATION WITH A SULFONYLUREA PLUS METFORMIN TO IMPROVE GLYCEMIC CONTROL IN PATIENTS WITH TYPE 2 DIABETES MELLITUS

U-692 USE OF VALSARTAN TO REDUCE CARDIOVASCULAR MORTALITY IN CLINICALLY STABLE PATIENTS WITH LEFT VENTRICULAR FAILURE OR LEFT VENTRICULAR DYSFUNCTION FOLLOWING MYOCARDIAL INFARCTION

U-693 THE RECOMMENDED INITIAL DOSE OF EQUETRO IS 400MG/DAY GIVEN IN DIVIDED DOSES, TWICE DAILY. THE DOSE SHOULD BE ADJUSTED IN 200MG DAILY INCREMENTS TO ACHIEVE OPTIMAL CLINICAL RESPONSE.

U-694 LENALIDOMIDE IS AN ANALOGUE OF THALIDOMIDE. THALIDOMIDE IS A KNOWN HUMAN TERATOGEN THAT CAUSES SEVERE LIFE-THREATENING HUMAN BIRTH DEFECTS. IF LENALIDOMIDE IS TAKEN DURING PREGNANCY, IT MAY CAUSE BIRTH DEFECTS OR DEATH TO AN UNBORN BABY.

U-695 TREATMENT OF PATIENTS WITH T-CELL ACUTE LYMPHOBLASTIC LEUKEMIA AND T-CELL

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

LYMPHOBLASTIC LYMPHOMA WHOSE DISEASE HAS NOT RESPONDED TO OR HAS RELAPSED FOLLOWING TREATMENT WITH AT LEAST TWO CHEMOTHERAPY REGIMENS

U-696 TREATMENT OF PATIENTS WITH T-CELL LYMPHOBLASTIC LYMPHOMA WHOSE DISEASE HAS NOT RESPONDED TO OR HAS RELAPSED FOLLOWING TREATMENT WITH AT LEAST TWO CHEMOTHERAPY REGIMENS

U-697 A METHOD OF USING RINFABATE RECOMBINANT (RHIGFBP-3) WITH MECASERMIN RECOMBINANT (RHIGF-1) TO PROMOTE LINEAR GROWTH IN THE TREATMENT OF PRIMARY IGF-1 DEFICIENCY

U-698 METHOD OF USING ANTAGONIST OF ARGININE VASOPRESSIN (AVA) V1A AND V2 RECEPTORS FOR INTRAVENOUS TREATMENT OF PATIENTS WITH EUVOLEMIC HYPONATREMIA

U-699 NASAL TREATMENT OF SEASONAL AND PERENNIAL ALLERGIC RHINITIS SYMPTOMS

U-700 TREATMENT AND PREVENTION OF OSTEOPOROSIS IN POSTMENOPAUSAL WOMEN

U-701 TREATMENT OF HYPERCHOLESTEROLEMIA AND/OR HYPERTRIGLYCERIDEMIA

U-702 TOPICAL AEROSOL HAIR REGROWTH TREATMENT

U-703 TREATMENT OF PROTEIN KINASE RELATED DISORDERS, SUCH AS GASTROINTESTINAL STROMAL TUMOR AND RENAL CELL CARCINOMA WITH SUNITINIB

U-704 METHOD OF ADMINISTERING INSULIN VIA INHALATION

U-705 TREATING CHRONIC ANGINA BY ADMINISTERING AN EXTENDED RELEASE FORM OF RANOLAZINE

U-706 TREATMENT OF BENIGN PROSTATIC HYPERPLASIA

U-707 ALLERGIC RHINITIS

U-708 TREATMENT OF CHRONIC NON-INFECTIOUS UVEITIS AFFECTING THE POSTERIOR SEGMENT OF THE EYE

U-709 METHOD OF COMBATING BACTERIA IN A PATIENT

U-710 A METHOD OF TREATING RESPIRATORY DISORDERS, E.G., ASTHMA, WHICH COMPRISES ADMINISTRATION BY INHALATION OF AN EFFECTIVE AMOUNT OF A PHARMACEUTICAL FORMULATION AS CLAIMED IN US PATENT NO. 5658549

U-711 ACUTE AND LONGER-TERM TREATMENT OF MAJOR DEPRESSIVE DISORDER

U-712 A METHOD OF USING A NICOTINIC ACID FORMULATION TO REDUCE ELEVATED TC, LDL-C AND TG LEVELS, AND RAISE HDL-C LEVELS IN PATIENTS WITH HYPERLIPIDEMIA

U-713 TREATMENT OF MILD TO MODERATE DEMENTIA OF THE ALZHEIMER'S TYPE

U-714 TOPICAL TREATMENT OF INTERDIGITAL TINEA PEDIS AND TINEA CORPORIS DUE TO TRICHOPHYTON RUBRUM, TRICHOPHYTON MENTAGROPHYTES OR EPIDERMOPHYTON FLOCCOSUM

U-715 FOR CLEANSING THE BOWEL IN PREPARATION FOR COLONOSCOPY, IN ADULTS 18 YEARS OF AGE OR OLDER

U-716 THE TREATMENT OR PREVENTION OF BRONCHOSPASM IN ADULTS AND CHILDREN 4 YEARS OF AGE AND OLDER WITH REVERSIBLE OBSTRUCTIVE AIRWAYS DISEASE AND THE PREVENTION OF EXERCISED-INDUCED BRONCHOSPASM IN PATIENTS 4 YEARS OF AGE AND OLDER

U-717 METHOD OF RELIEVING OR PREVENTING CONSTIPATION IN A HUMAN CONSTIPATED PATIENT

U-718 TREATMENT OF FUNGAL INFECTIONS

U-719 TREATMENT OF PSYCHOSIS

U-720 TREATMENT OF NEUROLEPTIC DISEASES

U-721 TREATMENT OF INFLUENZA

U-722 PROPHYLAXIS OF INFLUENZA

U-723 PROPHYLACTIC TREATMENT OF MIGRAINE

U-724 METHOD OF TREATING SEIZURES

U-725 ALLERGIC RHINITIS AND URTICARIA

U-726 ALLERGIC RHINITIS

U-727 FOR THE TREATMENT OF ATTENTION DEFICIT HYPERACTIVITY DISORDER (ADHD)

U-728 METHOD FOR TREATING BACTERIAL INFECTION

U-729 TREATMENT OF GASTROESOPHAGEAL REFLUX DISEASE (GERD), RISK-REDUCTION OF NSAID-ASSOCIATED GASTRIC ULCER, H. PYLORI ERADICATION TO REDUCE THE RISK OF DUODENAL ULCER RECURRENCE

U-730 USE AS A NASAL SPRAY FOR TREATMENT OF THE SYMPTOMS OF SEASONAL ALLERGIC RHINITIS

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

AND VASOMOTOR RHINITIS

- U-731 USE IN COMBINATION WITH DEXAMETHASONE IS INDICATED FOR THE TREATMENT OF PATIENTS WITH NEWLY DIAGNOSED MULTIPLE MYELOMA
- U-732 ACUTE TREATMENT OF THE CUTANEOUS MANIFESTATIONS OF MODERATE TO SEVERE ERYTHEMA NODOSUM LEPROSUM (ENL)
- U-733 MAINTENANCE THERAPY FOR PREVENTION AND SUPPRESSION OF THE CUTANEOUS MANIFESTATIONS OF ENL RECURRENCE
- U-734 FIRST LINE THERAPY FOR TYPE 2 DIABETES MELLITUS
- U-735 METHOD OF TREATING CHRONIC IRON OVERLOAD
- U-736 METHOD FOR IONTOPHORETIC TRANSDERMAL DELIVERY OF FENTANYL HYDROCHLORIDE
- U-737 DISINFECTION OF PATIENT SKIN PRIOR TO AN INVASIVE PROCEDURE
- U-738 INDICATED FOR THE LONG-TERM, TWICE-DAILY MAINTENANCE TREATMENT OF ASTHMA IN PATIENTS 12 YEARS OF AGE OR OLDER
- U-739 METHOD FOR TREATING CONSTIPATION BY OPENING CIC CHANNELS IN A MAMMALIAN SUBJECT
- U-740 FOR THE TREATMENT OF PATIENTS WITH PRIMARY BILIARY CIRRHOSIS
- U-741 COMBINATION THERAPY WITH CISPLATIN FOR THE TREATMENT OF LATE STAGE CERVICAL CANCER
- U-742 TWICE DAILY TOPICAL TREATMENT OF MODERATE TO SEVERE PLAQUE PSORIASIS.
- U-743 ONCE A DAY TOPICAL TREATMENT OF THE INFLAMMATORY LESIONS OF ROSACEA
- U-744 TREATMENT OF HIV INFECTION IN ANTIRETROVIRAL TREATMENT-EXPERIENCED ADULT PATIENTS
- U-745 TREATMENT OR PREVENTION OF EMESIS
- U-746 PREVENTION OR TREATMENT OF NAUSEA OR EMESIS INDUCED BY A CANCER CHEMOTHERAPEUTIC AGENT
- U-747 PREVENTION OR TREATMENT OF POST-OPERATIVE NAUSEA AND VOMITING
- U-748 A METHOD FOR THE TREATMENT OF A PROTEIN TYROSINE KINASE-ASSOCIATED DISORDER
- U-749 METHOD OF CONTRACEPTION
- U-750 TREATMENT OF HIV-1 INFECTION IN ADULTS
- U-751 ONCE DAILY DOSING OF BUDESONIDE VIA NEBULIZER FOR THE TREATMENT OF ASTHMA
- U-752 SUNSCREEN
- U-753 AS AN ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN PATIENTS WITH TYPE 2 DIABETES
- U-754 USE FOR THE LONG-TERM MAINTENANCE TREATMENT OF ASTHMA
- U-755 TREATMENT OF ANOREXIA, CACHEXIA, OR AN UNEXPLAINED, SIGNIFICANT WEIGHT LOSS IN PATIENTS WITH A DIAGNOSIS OF ACQUIRED IMMUNODEFICIENCY SYNDROME (AIDS)
- U-756 ADDITION OF ONCE-WEEKLY DOSING FOR THE TREATMENT TO INCREASE BONE MASS IN MEN WITH OSTEOPOROSIS
- U-757 USE AS A BILE ACID SEQUESTRANT FOR LOWERING CHOLESTEROL
- U-758 TREATMENT OF SYMPTOMS OF PREMENSTRUAL DYSPHORIC DISORDER
- U-759 METHOD OF USE OF ADMINISTERING LEVOTHYROXINE
- U-760 PROPHYLAXIS OF INVASIVE ASPERGILLUS AND CANDIDA INFECTIONS AND TREATMENT OF OROPHARYNGEAL CANDIDIASIS
- U-761 TREATMENT OF SCHIZOPHRENIA INCLUDING MAINTAINING STABILITY IN PATIENTS WITH SCHIZOPHRENIA
- U-762 TREATMENT OF CHRONIC OBSTRUCTIVE PULMONARY DISEASE
- U-763 ADMINISTRATION OF ARIPIPIRAZOLE BY INJECTION
- U-764 TREATMENT OF SCHIZOPHRENIA
- U-765 METHOD OF TREATING ALLERGIC CONJUNCTIVITIS
- U-766 TREATMENT OF SEIZURES
- U-767 MANAGEMENT OF BREAKTHROUGH PAIN IN PATIENTS WITH CANCER

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

- U-768 A METHOD OF REDUCING THE CAPACITY OF EXTENDED RELEASE NICOTINIC ACID TO PROVOKE A FLUSHING REACTION BY PRETREATING AN INDIVIDUAL WITH A FLUSH INHIBITING AGENT PRIOR TO THE ADMINISTRATION OF THE EXTENDED RELEASE NICOTINIC ACID
- U-769 REVLIMID (LENALIDOMIDE) IN COMBINATION WITH DEXAMETHASONE IS INDICATED FOR THE TREATMENT OF MULTIPLE MYELOMA PATIENTS WHO HAVE RECEIVED AT LEAST ONE PRIOR THERAPY
- U-770 LONG-TERM TREATMENT OF PATHOLOGICAL HYPERSECRETORY CONDITIONS
- U-771 METHOD FOR THE TREATMENT OF DIABETES MELLITUS, SUCH AS TYPE 1 DIABETES MELLITUS OR TYPE 2 DIABETES MELLITUS, IN A HUMAN PATIENT
- U-772 RELIEF OF SYMPTOMS ASSOCIATED WITH SEASONAL ALLERGIC RHINITIS IN CHILDREN 2 TO 11 YEARS AND FOR THE RELIEF OF SYMPTOMS ASSOCIATED WITH UNCOMPLICATED SKIN MANIFESTATIONS OF CHRONIC IDIOPATHIC URTICARIA IN CHILDREN 6 MONTHS TO 11 YEARS
- U-773 PATHOLOGICAL HYPERSECRETORY CONDITIONS
- U-774 METHOD OF TREATING TYPE 2 DIABETES MELLITUS BY ADMINISTERING A DIPEPTIDYL PEPTIDASE-IV INHIBITOR
- U-775 METHOD OF TREATING TYPE 2 DIABETES MELLITUS BY ADMINISTERING A DIPEPTIDYL PEPTIDASE-IV INHIBITOR IN COMBINATION WITH METFORMIN AND/OR A SULFONYLUREA
- U-776 TREATMENT OF CUTANEOUS MANIFESTATION IN PATIENTS WITH CUTANEOUS T-CELL LYMPHOMA (CTCL) WHO HAVE PROGRESSIVE, PERSISTENT OR RECURRENT DISEASE ON OR FOLLOWING TWO SYSTEMIC THERAPIES.
- U-777 DECREASING MORTALITY CAUSED BY CONGESTIVE HEART FAILURE
- U-778 REDUCTION OF ELEVATED INTRAOCULAR PRESSURE IN PATIENTS WITH OPEN ANGLE GLAUCOMA OR OCULAR HYPERTENSION
- U-779 A METHOD FOR TREATMENT OF A CANCER, WHEREIN THE CANCER IS CHRONIC MYELOGENOUS LEUKEMIA
- U-780 A METHOD FOR THE TREATMENT OF CANCER
- U-781 FOR TREATMENT OF ADULT PATIENTS WITH TYPE 2 DIABETES MELLITUS WHO ARE NAIVE TO PHARMACOLOGIC THERAPY
- U-782 TREATMENT OF CHRONIC HEPATITIS B IN ADULT PATIENTS WITH EVIDENCE OF VIRAL REPLICATION AND EITHER EVIDENCE OF PERSISTENT ELEVATIONS IN SERUM AMINOTRANSFERASES (ALT OR AST) OR HISTOLOGICALLY ACTIVE DISEASE
- U-783 DESONATE GEL IS INDICATED FOR THE TREATMENT OF MILD TO MODERATE ATOPIC DERMATITIS IN PATIENTS 3 MONTHS OF AGE AND OLDER
- U-784 TREATMENT OF MODERATE TO SEVERE PRIMARY RESTLESS LEGS SYNDROME (RLS)
- U-785 USE AS REPLACEMENT SOLUTION, HEMOFILTRATION SOLUTION OR HEMODIAFILTRATION SOLUTION IN CONTINUOUS RENAL REPLACEMENT THERAPY
- U-786 PRODUCT IS APPROVED FOR THE TOPICAL TREATMENT OF TINEA PEDIS
- U-787 MAINTENANCE TREATMENT OF ASTHMA AS PROPHYLACTIC THERAPY IN ADULT AND PEDIATRIC PATIENTS SIX YEARS OF AGE OR OLDER, INCLUDING PATIENTS REQUIRING ORAL CORTICOSTEROID THERAPY FOR ASTHMA
- U-788 METHOD OF TREATING PSYCHIATRIC SYMPTOMS ASSOCIATED WITH PREMENSTRUAL DISORDERS USING PAROXETINE
- U-789 TREATMENT OF KNOWN OR SUSPECTED CYANIDE POISONING
- U-790 FORTEO IS INDICATED FOR THE TREATMENT OF POST MENOPAUSAL WOMEN WITH OSTEOPOROSIS WHO ARE AT RISK FOR FRACTURE. FORTEO CAN BE USED BY PEOPLE WHO HAVE HAD A FRACTURE RELATED TO OSTEOPOROSIS
- U-791 GLEEVEC IS ALSO INDICATED FOR THE TREATMENT OF PATIENTS WITH KIT (CD117) POSITIVE UNRESECTABLE AND/OR METASTATIC MALIGNANT GASTROINTESTINAL STROMAL TUMORS (GIST)
- U-792 TREATMENT OF SEBORRHEA DERMATITIS IN HUMANS
- U-793 FOR THE LONG TERM TREATMENT, TWICE DAILY (MORNING AND EVENING) MAINTENANCE TREATMENT OF BRONCHOCONSTRICTION IN PATIENTS WITH CHRONIC OBSTRUCTIVE PULMONARY DISEASE (COPD), INCLUDING CHRONIC BRONCHITIS AND EMPHYSEMA
- U-794 CLOSURE OF A CLINICALLY SIGNIFICANT PATENT DUCTUS ARTERIOSUS IN PREMATURE INFANTS WEIGHING BETWEEN 500 AND 1500G, WHO ARE NO MORE THAN 32 WEEKS GESTATIONAL AGE WHEN USUAL MEDICAL MANAGEMENT IS INEFFECTIVE
- U-795 METHOD FOR INHIBITING NOREPINEPHRINE UPTAKE

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-796 METHOD OF TREATING DEPRESSION

U-797 METHOD OF TREATING ANXIETY

U-798 TREATMENT AND PREVENTION OF OSTEOPOROSIS IN POSTMENOPAUSAL WOMEN BY ONCE-MONTHLY ORAL ADMINISTRATION OF IBANDRONATE SODIUM MONOHYDRATE EQUIVALENT TO 150MG OF IBANDRONIC ACID

U-799 METHOD FOR INHIBITING SEROTONIN UPTAKE

U-800 TREATMENT OF PATIENTS WITH ADVANCED OR METASTATIC BREAST CANCER WHOSE TUMORS OVEREXPRESS HER2 AND WHO HAVE RECEIVED PRIOR THERAPY INCLUDING ANTHRACYCLINE, A TAXANE AND TRASTUZUMAB

U-801 METHOD OF TREATING CANCER

U-802 METHOD OF TREATING TYPE 2 DIABETES MELLITUS BY ADMINISTERING A DIPEPTIDYL PEPTIDASE-IV INHIBITOR

U-803 METHOD OF TREATING TYPE 2 DIABETES MELLITUS BY ADMINISTERING A DIPEPTIDYL PEPTIDASE-IV INHIBITOR IN COMBINATION WITH METFORMIN

U-804 TREATMENT OF ACTINIC KERATOSES BY PHOTODYNAMIC THERAPY

U-805 TREATMENT OF IMPETIGO DUE TO STAPHYLOCOCCUS AUREUS OR STREPTOCOCCUS PYOGENES

U-806 INTRATHECAL TREATMENT OF LYMPHOMATOUS MENINGITIS

U-807 PREVENTION OF EXERCISE-INDUCED BRONCHOCONSTRICTION

U-808 THE TREATMENT OF THE SYMPTOMS OF SEASONAL AND PERENNIAL ALLERGIC RHINITIS IN PATIENTS 2 YEARS OF AGE AND OLDER

U-809 TREATMENT OF CHRONIC IDIOPATHIC URTICARIA

U-810 METHOD OF TREATMENT TO ALLEVIATE INFLAMMATION OF THE EYE

U-811 RELIEF OF SYMPTOMS ASSOCIATED WITH SEASONAL AND PERENNIAL ALLERGIC RHINITIS AND TREATMENT OF THE UNCOMPLICATED SKIN MANIFESTATIONS OF CHRONIC IDIOPATHIC URTICARIA

U-812 RELIEF OF SYMPTOMS ASSOCIATED WITH SEASONAL AND PERENNIAL ALLERGIC RHINITIS

U-813 MAINTENANCE TREATMENT OF BRONCHOCONSTRICTION IN PATIENTS WITH CHRONIC OBSTRUCTIVE PULMONARY DISEASE (COPD)

U-814 TREATMENT OF SCHIZOPHRENIA

U-815 TREATS COLD SORES/FEVER BLISTERS ON THE FACE OR LIPS. SHORTENS HEALING TIME AND DURATION OF SYMPTOMS: TINGLING, PAIN, BURNING AND/OR ITCHING

U-816 DEPRESSION, PANIC DISORDER, PREMENSTRUAL DISORDERS AND SOCIAL ANXIETY DISORDER

U-817 NASAL ADMINISTRATION OF CYANOCOBALAMIN

U-818 TOPICAL TREATMENT OF ACNE VULGARIS

U-819 MANAGEMENT OF FIBROMYALGIA

U-820 IMPROVED WAKEFULNESS IN PATIENTS WITH EXCESSIVE SLEEPINESS ASSOCIATED WITH NARCOLEPSY, OBSTRUCTIVE SLEEP APNEA/HYPOPNEA SYNDROME, AND SHIFT WORK SLEEP DISORDER

U-821 METHOD OF INHIBITING ENTHOTHELIN RECEPTORS BY ADMINISTERING AMBRISENTAN TO A PATIENT TO TREAT PULMONARY ARTERIAL HYPERTENSION.

U-822 USE IN LIPID MANAGEMENT

U-823 RELIEF OF SYMPTOMS ASSOCIATED WITH SEASONAL ALLERGIC RHINITIS AND FOR THE TREATMENT OF UNCOMPLICATED SKIN MANIFESTATIONS OF CHRONIC IDIOPATHIC URTICARIA IN CHILDREN 6 TO 11 YEARS OF AGE

U-824 METHOD OF TREATING PATIENTS INFECTED WITH CCR5-TROPIC HIV-1

U-825 USE FOR PREVENTION OF BREAST CANCER

U-826 RELIEF OF MODERATE TO SEVERE PAIN

U-827 USE FOR TREATMENT OF DIABETES, PARTICULARLY TYPE 2 DIABETES

U-828 PREVENTION OF PREGNANCY IN WOMEN WHO ELECT TO USE ORAL CONTRACEPTIVES AS A METHOD OF CONTRACEPTION

U-829 TREATMENT OF EXTRAVASATION RESULTING FROM IV ANTHRACYCLINE CHEMOTHERAPY

U-830 TREATMENT OF RELAPSED SMALL CELL LUNG CANCER

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-831 METHOD OF ADMINISTERING LANREOTIDE ACETATE

U-832 ZINGO IS INDICATED FOR THE USE ON INTACT SKIN TO PROVIDE LOCAL ANALGESIA PRIOR TO VENIPUNCTURE OR INTRAVENOUS CANNULATION.

U-833 METHOD OF TREATING PAIN USING A PHARMACEUTICALLY ACCEPTABLE SALT OF ROPIVACAINE AND ADMINISTERING A COMPOSITION CONTAINING LESS THAN 0.25% BY WEIGHT OF ROPIVACAINE

U-834 INVIRASE IN COMBINATION WITH RITONAVIR AND OTHER ANTIRETROVIRAL AGENTS IS INDICATED FOR THE TREATMENT OF HIV INFECTION

U-835 RELIEF OF THE INFLAMMATORY AND PRURITIC MANIFESTATIONS OF ATOPIC DERMATITIS IN PATIENTS ONE YEAR OF AGE OR OLDER

U-836 A METHOD FOR THE TREATMENT OF LEUKEMIAS

U-837 GASTROINTESTINAL LAVAGE INDICATED FOR CLEANSING OF THE COLON AS A PREPARATION FOR COLONOSCOPY IN ADULTS

U-838 METHOD OF TREATING PAIN USING A PHARMACEUTICALLY ACCEPTABLE SALT OF ROPIVACAINE AND ADMINISTERING A COMPOSITION CONTAINING LESS THAN 0.5% BY WEIGHT OF ROPIVACAINE

U-839 TREATMENT OF MAJOR DEPRESSIVE DISORDER (MDD)

U-840 TREATMENT FOR TYPE 2 DIABETES MELLITUS

U-841 INDICATED FOR THE LONG-TERM, MAINTENANCE TREATMENT OF ASTHMA IN PATIENTS 12 YEARS OF AGE AND OLDER

U-842 INDICATED FOR THE TREATMENT OF ATTENTION-DEFICIT/HYPERACTIVITY DISORDER (ADHD)

U-843 METHOD FOR ADMINISTRATION OF TESTOSTERONE

U-844 PREFEST IS INDICATED IN WOMEN WHO HAVE A UTERUS FOR THE TREATMENT OF MODERATE TO SEVERE VASOMOTOR SYMPTOMS ASSOCIATED WITH MENOPAUSE; TREATMENT OF VULVAR AND VAGINAL ATROPHY; PREVENTION OF OSTEOPOROSIS

U-845 TREATMENT OF PATIENTS WITH CANDIDEMIA, ACUTE DISSEMINATED CANDIDIASIS, CANDIDA PERITONITIS AND ABSCESES

U-846 USE FOR DELINEATION (VISUALIZATION) DURING A VITRECTOMY SURGICAL PROCEDURE

U-847 ADJUNCTIVE THERAPY TO DIET IN ADULTS TO REDUCE LDL-C, TRIGLYCERIDES AND APO B, AND INCREASE HDL-C IN PATIENTS WITH PRIMARY HYPERCHOLESTEROLEMIA OR MIXED DYSLIPIDEMIA (TYPES IIA, IIB) AND TO TREAT HYPERTRIGLYCERIDEMIA (TYPES IV, V)

U-848 ACUTE TREATMENT OF MIGRAINE WITH OR WITHOUT AURA

U-849 REDUCTION OF ELEVATED INTRAOCULAR PRESSURE (IOP) IN PATIENTS WITH GLAUCOMA OR OCULAR HYPERTENSION WHO REQUIRE ADJUNCTIVE OR REPLACEMENT THERAPY DUE TO INADEQUATELY CONTROLLED IOP. DOSE IS ONE DROP OF COMBIGAN IN THE AFFECTED EYE TWICE DAILY

U-850 PREVENTION OR TREATMENT OF NAUSEA OR EMESIS INDUCED BY A CANCER CHEMOTHERAPEUTIC AGENT

U-851 TREATMENT OF TYPE 2 DIABETES MELLITUS

U-852 RELIEF OF SYMPTOMS ASSOCIATED WITH SEASONAL AND PERENNIAL ALLERGIC RHINITIS

U-853 TREATMENT OR PREVENTION OF EMESIS

U-854 PREVENTION OF CMV DISEASE IN KIDNEY, HEART, AND KIDNEY-PANCREAS TRANSPLANT PATIENTS AT HIGH RISK (DONOR CMV SEROPOSITIVE/RECIPIENT CMV SERONEGATIVE)

U-855 METHOD TO INDUCE NATRIURESIS, DIURESIS AND/OR VASODILATION

U-856 SUPPORT EMBRYO IMPLANTATION AND EARLY PREGNANCY BY SUPPLEMENTATION OF CORPUS LUTEAL FUNCTION AS PART OF AN ASSISTED REPRODUCTIVE TECHNOLOGY (ART) TREATMENT PROGRAM FOR INFERTILE WOMEN

U-857 INHIBITION OF TRANSPLANT REJECTION

U-858 PEDIATRIC USE AGED 1-11 YEARS, GERD AND EROSIIVE ESOPHAGITIS

U-859 EROSIIVE ESOPHAGITIS, HYPERSECRETORY CONDITIONS INCLUDING ZOLLINGER-ELLISON SYNDROME, MAINTENANCE OF HEALING OF EROSIIVE ESOPHAGITIS AND REDUCTION OF SYMPTOMS IN PATIENTS WITH GERD

U-860 FOR THE APPROVED USES AND CONDITIONS OF USE, INCLUDING DEPRESSION

U-861 RELIEF OF THE INFLAMMATORY AND PRURITIC MANIFESTATIONS OF CORTICOSTEROID RESPONSIVE DERMATOSES IN PATIENTS 12 YEARS OF AGE OR OLDER

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

- U-862 ADJUNCT TO DIET TO REDUCE ELEVATED TOTAL-C, LDL-C, NON-HDL-C, APO B, TG, AND LP(A) LEVELS AND TO INCREASE HDL-C IN PATIENTS WITH PRIMARY HYPERCHOLESTEROLEMIA, MIXED DYSLIPIDEMIA, AND HYPERTRIGLYCERIDEMIA
- U-863 TAKING ASPIRIN OR NON-STEROIDAL ANTI-INFLAMMATORY MEDICATIONS APPROXIMATELY 30 MINUTES BEFORE DOSING CAN MINIMIZE FLUSHING, A COMMON SIDE EFFECT OF NIACIN THERAPY
- U-864 PEDIATRIC USE AGES 1-2 YEARS, GERD AND EROSIIVE ESOPHAGITIS
- U-865 TREATMENT OF A WOMAN WITH OSTEOPOROSIS AND A HIGH RISK FOR BONE FRACTURE BY REDUCING THE RISK OF VERTEBRAL AND NONVERTEBRAL BONE FRACTURE
- U-866 THE LABEL REFERENCES THE EFFECTS OF THE ACTIVE INGREDIENT OF REVLIMID UPON CYTOKINES
- U-867 TREATMENT OF MIGRAINE
- U-868 METHOD OF USING ANTAGONIST OF ARGININE VASOPRESSIN (AVA) V1A AND V2 RECEPTORS FOR INTRAVENOUS TREATMENT OF PATIENTS WITH HYPERVOLEMIC HYONATREMIA
- U-869 METHOD FOR STIMULATING CORONARY VASODILATION FOR PURPOSES OF IMAGING THE HEART
- U-870 METHOD OF PRODUCING CORONARY VASODILATION WITHOUT PERIPHERAL VASODILATION
- U-871 METHOD OF REDUCING RISK OF MYOCARDIAL INFARCTION, STROKE AND DEATH
- U-872 TWICE DAILY MAINTENANCE TREATMENT OF AIRFLOW OBSTRUCTION IN PATIENTS WITH CHRONIC OBSTRUCTIVE PULMONARY DISEASE (COPD), INCLUDING CHRONIC BRONCHITIS AND EMPHYSEMA. TO REDUCE EXACERBATIONS OF COPD IN PATIENTS WITH A HISTORY OF EXACERBATIONS
- U-873 METHOD OF TREATING CONSTIPATION IN A PATIENT WITH IRRITABLE BOWEL SYNDROME BY OPENING CHLORIDE CHANNELS (CIC)
- U-874 METHOD OF TREATING CONSTIPATION IN A PATIENT WITH IRRITABLE BOWEL SYNDROME
- U-875 FIRST-LINE TREATMENT OF LOCALLY ADVANCED UNRESECTABLE OR METASTATIC PANCREATIC CANCER, IN COMBINATION WITH GEMCITABINE
- U-876 TREATMENT OF MIGRAINE WITH OR WITHOUT AURA
- U-877 FOR USE AS ADJUNCTIVE THERAPY IN THE TREATMENT OF PEPTIC ULCER
- U-878 A METHOD FOR BINDING A PERIPHERAL OPIOID RECEPTOR
- U-879 A METHOD OF TREATING OR PREVENTING ILEUS
- U-880 ENDOMETRIN IS A PROGESTERONE INDICATED TO SUPPORT EMBRYO IMPLANTATION AND EARLY PREGNANCY BY SUPPLEMENTATION OF CORPUS LUTEAL FUNCTION AS PART OF AN ASSISTED REPRODUCTIVE TECHNOLOGY (ART) TREATMENT PROGRAM FOR INFERTILE WOMEN
- U-881 TREATMENT OF NON-SMALL CELL LUNG CANCER
- U-882 MANAGEMENT OF FIBROMYALGIA (FM)
- U-883 TREATMENT OF GASTROINTESTINAL STROMAL TUMOR WITH SUNITINIB
- U-884 TREATMENT OF PATIENTS WITH MULTIPLE MYELOMA
- U-885 TREATMENT OF PATIENTS WITH MANTLE CELL LYMPHOMA WHO HAVE RECEIVED AT LEAST 1 PRIOR THERAPY
- U-886 ADMINISTERING DESLORATADINE TO TREAT THE SYMPTOMS OF PERENNIAL ALLERGIC RHINITIS, SEASONAL ALLERGIC RHINITIS, OR CHRONIC IDIOPATHIC URTICARIA
- U-887 TREATMENT AND PREVENTION OF OSTEOPOROSIS
- U-888 FEMALE HORMONE REPLACEMENT THERAPY FOR POSTMENOPAUSAL WOMEN
- U-889 MENOPAUSAL AND POSTMENOPAUSAL DISORDERS (INCLUDING VASOMOTOR SYMPTOMS ASSOCIATED WITH MENOPAUSE)
- U-890 REDUCTION OF SERUM PHOSPHATE IN PATIENTS WITH END STAGE RENAL DISEASE
- U-891 USE AS AN INTRAOCULAR IRRIGATING SOLUTION DURING SURGICAL PROCEDURES INVOLVING PERFUSION OF THE EYE
- U-892 TREATMENT OF CUTANEOUS MANIFESTATIONS IN PATIENTS WITH CUTANEOUS T-CELL LYMPHOMA (CTCL)
- U-893 CLEVIPREX IS A DIHYDROPYRIDINE CALCIUM CHANNEL BLOCKER INDICATED FOR THE REDUCTION OF BLOOD PRESSURE WHEN ORAL THERAPY IS NOT FEASIBLE OR NOT DESIRABLE
- U-894 TREATMENT OF COLD SORES IN PEDIATRIC PATIENTS TWELVE YEARS OF AGE AND OLDER

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-895 TREATMENT OF HIV INFECTION IN COMBINATION WITH OTHER ANTIRETROVIRAL AGENTS

U-896 TREATMENT OF NASAL SYMPTOMS OF SEASONAL AND PERENNIAL ALLERGIC RHINITIS IN ADULTS AND CHILDREN TWO YEARS OF AGE AND OLDER

U-897 METHOD OF TREATING TONSILLITIS AND/OR PHARYNGITIS SECONDARY TO STREPTOCOCCUS PYOGENES IN A ONCE-A-DAY AMOXICILLIN PRODUCT

U-898 USE OF GLUTAMINE TOGETHER WITH GROWTH HORMONE FOR THE TREATMENT OF PATIENTS WITH SHORT BOWEL SYNDROME

U-899 USE OF THALIDOMIDE IN COMBINATION WITH DEXAMETHASONE FOR THE TREATMENT OF PATIENTS WITH NEWLY DIAGNOSED MULTIPLE MYELOMA

U-900 INTEGRASE INHIBITION FOR THE TREATMENT OF HIV INFECTION

U-901 PREVENTION OF POSTOPERATIVE NAUSEA AND VOMITING

U-902 USE IN THE TREATMENT OF THE SIGNS AND SYMPTOMS OF BENIGN PROSTATIC HYPERPLASIA (BPH)

U-903 TREATMENT OF HUMAN IMMUNODEFICIENCY VIRUS (HIV) IN ADULT PATIENTS

U-904 TREATMENT OF MODERATE TO SEVERE VASOMOTOR SYMPTOMS ASSOCIATED WITH MENOPAUSE

U-905 TREATMENT OF MODERATE TO SEVERE VAGINAL DRYNESS AND PAIN WITH INTERCOURSE, SYMPTOMS OF VULVAR AND VAGINAL ATROPHY, ASSOCIATED WITH MENOPAUSE

U-906 PROPHYLAXIS OF ORGAN REJECTION IN KIDNEY, LIVER AND HEART ALLOGENIC TRANSPLANTS; TREATMENT OF PATIENTS WITH SEVERE ACTIVE, RHEUMATOID ARTHRITIS; TREATMENT OF ADULT, NONIMMUNOCOMPROMISED PATIENTS WITH SEVERE, RECALCITRANT, PLAQUE PSORIASIS

U-907 FOR THE MAINTENANCE OF REMISSION OF ULCERATIVE COLITIS IN SUBJECTS 18 YEARS OF AGE AND OLDER

U-908 PROPHYLAXIS OF ORGAN REJECTION IN PATIENTS RECEIVING ALLOGENEIC RENAL TRANSPLANTS

U-909 TREATMENT OF CYSTIC FIBROSIS PATIENTS WITH PSEUDOMONAS AERUGINOSA

U-910 TREATMENT OF METASTATIC CARCINOMA OF THE OVARY AFTER FAILURE OF INITIAL OR SUBSEQUENT CHEMOTHERAPY

U-911 METHOD OF TREATING, AS ADJUNCTIVE THERAPY, PARTIAL-ONSET SEIZURES IN A PATIENT WITH EPILEPSY AGED 17 YEARS AND OLDER WHEN ORAL TREATMENT IS TEMPORARILY NOT FEASIBLE

U-912 SEDATION OF NON-INTUBATED PATIENTS PRIOR TO AND/OR DURING SURGICAL AND OTHER PROCEDURES

U-913 TREATMENT OF OVERACTIVE BLADDER WITH SYMPTOMS OF URGE URINARY INCONTINENCE, URGENCY, AND FREQUENCY

U-914 METHOD OF TREATING, AS ADJUNCTIVE THERAPY, PARTIAL-ONSET SEIZURES IN A PATIENT WITH EPILEPSY AGED 17 YEARS AND OLDER

U-915 TREATMENT OF MUSCULOSKELETAL CONDITIONS

U-916 TOPICAL TREATMENT OF ACNE VULGARIS IN PATIENTS 12 YEARS OR OLDER

U-917 TREATMENT OF INFLAMMATORY LESIONS OF NON-NODULAR MODERATE TO SEVERE ACNE VULGARIS

U-918 TO TREAT OR PREVENT INFECTIONS CAUSED BY SUSCEPTIBLE BACTERIA USING DELAYED-RELEASE TABLETS CONSISTING OF DOXYCYCLINE HYCLATE COATED PELLETS IN A TABLET

U-919 FOR THE TREATMENT OF DERMATITIS

U-920 STEROID-RESPONSIVE INFLAMMATORY OCULAR CONDITIONS FOR WHICH A CORTICOSTEROID IS INDICATED AND WHERE SUPERFICIAL BACTERIAL OCULAR INFECTION OR A RISK OF BACTERIAL OCULAR INFECTION EXISTS

U-921 TREATMENT OF ACNE VULGARIS

U-922 FOR THE TREATMENT OF FUNGAL INFECTIONS

U-923 METHOD OF TREATING OPHTHALMIC INFLAMMATION AND INFECTION

U-924 TREATMENT OF MILD TO MODERATE INFECTION CAUSED BY SUSCEPTIBLE STRAINS

U-925 TREATMENT OF ONLY INFLAMMATORY LESIONS (PAPULES AND PUSTULES) OF ROSACEA

U-926 MGT SPECIFIC BACTERIAL INFECTIONS. TREATMENT PTS W/ COMMUNITY ACQUIRED PNEUMONIA OR BACTERIAL SINUSITIS DUE TO CONFIRMED, OR SUSPECTED B-LACTAMASE PRODUCING PATHOGENS & S. PNEUMONIAE WITH REDUCED SUSCEPTIBILITY TO PENICILLIN (MIC=2MC/ML)

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-927 METHOD FOR INCREASING TEAR PRODUCTION

U-928 TREATMENT OF BACTERIAL INFECTIOUS DISEASE

U-929 TREATMENT OF OBSESSIVE COMPULSIVE DISORDER TREATABLE WITH AN SSRI

U-930 TREATMENT OF IDIOPATHIC THROMBOCYTOPENIC PURPURA (ITP)

U-931 RELIEF OF MODERATE TO SEVERE ACUTE PAIN

U-932 PYLERA CAPSULES, IN COMBINATION WITH OMEPRAZOLE ARE INDICATED FOR THE TREATMENT OF PATIENTS WITH HELICOBACTER PYLORI INFECTION AND DUODENAL ULCER DISEASE TO ERADICATE H. PYLORI

U-933 FOR THE TREATMENT OF PATIENTS WITH HELICOBACTER PYLORI INFECTION AND DUODENAL ULCER DISEASE TO ERADICATE H. PYLORI. THE ERADICATION OF HELICOBACTER PYLORI HAS BEEN SHOWN TO REDUCE THE RISK OF DUODENAL ULCER RECURRENCE

U-934 IN COMBINATION WITH GRANULOCYTE-COLONY STIMULATING FACTOR (G-CSF) TO MOBILIZE HEMATOPOIETIC STEM CELL TO THE PERIPHERAL BLOOD FOR COLLECTION AND SUBSEQUENT AUTOLOGOUS TRANSPLANTATION WITH NON-HODGKINS LYMPHOMA AND MULTIPLE MYELOMA

U-935 TREATMENT OF HUMAN IMMUNODEFICIENCY VIRUS (HIV) INFECTION IN ADULT PATIENTS, AND TREATMENT OF HUMAN IMMUNODEFICIENCY VIRUS (HIV) IN PEDIATRIC PATIENTS 6 YEARS OF AGE AND OLDER

U-936 USE IN COMBINATION WITH GRANULOCYTE-COLONY STIMULATING FACTOR (G-CSF) TO MOBILIZE HEMATOPOIETIC STEM CELLS TO PERIPHERAL BLOOD FOR COLLECTION & SUBSEQUENT AUTOLOGOUS TRANSPLANTATION IN PATIENTS WITH NON-HODGKIN'S LYMPHOMA & MULTIPLE MYELOMA

U-937 TREATMENT OF PROSTATE CANCER

U-938 TREATMENT OF HAIR LOSS AND HYPOTRICHOSIS OF THE EYELASHES BY INCREASING THEIR GROWTH INCLUDING LENGTH, THICKNESS AND DARKNESS

U-939 TREATMENT OF HYPOTRICHOSIS OF THE EYELASHES BY INCREASING AND STIMULATING THEIR GROWTH INCLUDING LENGTH, THICKNESS AND DARKNESS

U-940 METHOD TO TREAT AIDS-RELATED KAPOSI'S SARCOMA

U-941 METHOD TO TREAT OVARIAN CANCER

U-942 METHOD TO TREAT MULTIPLE MYELOMA

U-943 GNRH ANTAGONIST INDICATED FOR TREATMENT OF PATIENTS WITH ADVANCED PROSTATE CANCER

U-944 TREATMENT OF PATIENTS WITH B-CELL CHRONIC LYMPHOCYTIC LEUKEMIA (CLL)

U-945 SEDATIVE-HYPNOTIC AGENT INDICATED FOR MONITORED ANESTHESIA CARE (MAC) SEDATION

U-946 TREATMENT OF BREAST CANCER

U-947 WHEN PATIENTS ARE UNABLE TO TAKE THE ORAL FORMULATIONS, PREVACID IV, FOR INJECTION IS INDICATED AS AN ALTERNATIVE FOR THE SHORT-TERM TREATMENT (UP TO 7 DAYS) OF ALL GRADES OF EROSIIVE ESOPHAGITIS

U-948 TREATMENT OF DIABETES MELLITUS

U-949 HEALING OF ALL GRADES OF EROSIIVE ESOPHAGITIS (EE) FOR UP TO 8 WEEKS

U-950 MAINTAIN HEALING OF EROSIIVE ESOPHAGITIS (EE) FOR UP TO 6 MONTHS

U-951 TREATMENT OF HEARTBURN ASSOCIATED WITH NON-EROSIVE GASTROESOPHAGEAL REFLUX DISEASE (GERD) FOR 4 WEEKS

U-952 USE AS AN ANALGESIC

U-953 METHOD OF TREATING OPHTHALMIC INFLAMMATION AND INFECTION

U-954 CHRONIC MANAGEMENT OF HYPERURICEMIA IN PATIENTS WITH GOUT. NOT RECOMMENDED FOR THE TREATMENT OF ASYMPTOMATIC HYPERURICEMIA

U-955 PROPHYLACTIC TREATMENT OF MIGRAINE

U-956 TREATMENT OF PATIENTS WITH H. PYLORI INFECTION AND DUODENAL ULCER DISEASE

U-957 A METHOD OF TREATING CANCER IN A PATIENT COMPRISING ADMINISTERING IXABEPILONE OR PHARMACEUTICAL COMPOSITIONS COMPRISING IXABEPILONE

U-958 METHOD OF TREATING PATIENT COMPRISING MIXING FIRST AND SECOND VIALS OF PRODUCT COMPRISING LYOPHILIZED IXABEPILONE TO PROVIDE AN EPOTHILONE ANALOG SOLUTION, DILUTING SOLUTION WITH A SUITABLE DILUENT TO PREPARE INTRAVENOUS FORMULATION FOR PT

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

- U-959 METHOD OF TREATING CANCER, IV ADMIN, LYOPHILIZED IXABEPILONE DILUTED, EVERY WEEK OR 3 WEEKS; LYOPHILIZED IXABEPILONE WITH SOLVENT (DEHYDRATED ETHANOL) DILUTED TO CONCENTRATION OF 0.1MG/ML TO 0.9MG/ML
- U-960 METHOD OF TREATING CANCER IN A PATIENT COMPRISING INTRAVENOUSLY ADMINISTERING TO THE PATIENT IXABEPILONE DILUTED IN A PARENTERAL DILUENT
- U-961 METHOD OF TREATING BREAST CANCER BY ADMINISTERING IXABEPILONE; A METHOD OF TREATING A CANCER RESPONSIBLE TO MICROTUBULE STABILIZATION BY ADMINISTERING IXABEPILONE
- U-962 SYMBYAX IS INDICATED FOR THE ACUTE TREATMENT OF TREATMENT RESISTANT DEPRESSION IN ADULTS
- U-963 PROZAC AND OLANZAPINE IN COMBINATION FOR THE ACUTE TREATMENT OF TREATMENT RESISTANT DEPRESSION IN ADULTS
- U-964 ZYPREXA ZYDIS AND FLUOXETINE IN COMBINATION FOR THE ACUTE TREATMENT OF TREATMENT RESISTANT DEPRESSION IN ADULTS
- U-965 USE OF IXABEPILONE IN COMBINATION WITH CAPECITABINE IN TREATMENT OF METASTASIS BREAST CANCER
- U-966 TREATMENT OF ASTHMA (MAINTENANCE AND PROPHYLACTIC THERAPY)
- U-967 A METHOD OF REVERSING SOFT-TISSUE ANESTHESIA I.E. ANESTHESIA OF THE LIP AND TONGUE, AND THE ASSOCIATED FUNCTIONAL DEFICITS RESULTING FROM AN INTRAORAL SUBMUCOSAL INJECTION OF A LOCAL ANESTHETIC
- U-968 A METHOD FOR IMPROVING GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS
- U-969 TREATMENT OF MIGRAINE
- U-970 TOPICAL TREATMENT OF LICE INFESTATIONS
- U-971 INDICATED FOR THE ACUTE TREATMENT OF ADULTS WITH SCHIZOPHRENIA
- U-972 MONOTHERAPY OR AS ADJUNCTIVE THERAPY TO LITHIUM OR VALPROATE FOR THE MAINTENANCE TREATMENT OF BIPOLAR I DISORDER
- U-973 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS WHO ARE ALREADY TREATED WITH PIOGLITAZONE AND METFORMIN OR WHO HAVE INADEQUATE GLYCEMIC CONTROL ON PIOGLITAZONE OR METFORMIN ALONE
- U-974 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN PATIENTS WITH TYPE 2 DIABETES WHO ARE ALREADY TREATED WITH A PIOGLITAZONE AND METFORMIN
- U-975 TREATMENT OF PULMONARY HYPERTENSION
- U-976 IMPROVEMENT OF GLYCEMIC CONTROL IN INDIVIDUALS WITH TYPE 2 DIABETES
- U-977 TREATMENT OF ACUTE, UNCOMPLICATED MALARIA INFECTION DUE TO PLASMODIUM FALCIPARUM IN PATIENTS OF 5KG BODYWEIGHT AND ABOVE
- U-978 METHOD OF TREATING HYPONATREMIA
- U-979 RELIEF OF MUSCLE SPASM
- U-980 NONSTEROIDAL ANTI-INFLAMMATORY DRUG INDICATED FOR RELIEF OF MILD TO MODERATE ACUTE PAIN
- U-981 MANAGEMENT OF MILD TO MODERATE PAIN, MANAGEMENT OF MODERATE TO SEVERE PAIN AS AN ADJUNCT TO OPIOID ANALGESICS, REDUCTION IN FEVER THROUGH ANTI-INFLAMMATORY, ANALGESIC, AND ANTIPYRETIC ACTIVITY
- U-982 A METHOD OF TREATING OSTEOPOROSIS
- U-983 METHOD OF TREATING OSTEOPOROSIS IN A POST-MENOPAUSAL WOMAN AT RISK FOR FRACTURE
- U-984 METHOD FOR THE TREATMENT OF A WOMAN WITH OSTEOPOROSIS AND AT RISK FOR BONE FRACTURE
- U-985 TREATMENT OF MACULAR EDEMA FOLLOWING BRANCH RETINAL VEIN OCCLUSION (BRVO) OR CENTRAL RETINAL VEIN OCCLUSION (CRVO)
- U-986 TREATMENT OF PATIENTS INFECTED WITH PEDICULUS HUMANUS CAPITIS (HEAD LICE AND THEIR OVA) OF THE SCALP HAIR
- U-987 TREATMENT OF SECONDARY HYPERPARATHYROIDISM IN PATIENTS WITH CHRONIC KIDNEY DISEASE ON DIALYSIS
- U-988 TREATMENT OF RHINITIS COMPRISING THE NASAL APPLICATION OF A PHARMACEUTICAL FORMULATION AS CLAIMED IN US PATENT 7541350
- U-989 FOR REDUCING BLOOD PHENYLALANINE LEVELS IN A HUMAN SUFFERING FROM HYPERPHENYLALANINEMIA

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-990 TREATMENT OF PROTOZOAL INFECTION

U-991 TREATMENT OR PROPHYLAXIS OF THROMBOSIS OR EMBOLISMS

U-992 REDUCTION OF THE RISK OF CARDIOVASCULAR HOSPITALIZATION

U-993 METHOD OF TREATING INFERTILITY

U-994 METHOD OF TREATMENT OF OSTEOPOROSIS WHEREIN THE OSTEOPOROSIS IS STEROID-INDUCED

U-995 METHOD FOR TREATING TYPE II DIABETES BY ADMINISTERING SAXAGLIPTIN

U-996 AN ADJUNCTIVE THERAPY TO DIET TO REDUCE ELEVATED TOTAL CHOLESTEROL (TC), LOW-DENSITY LIPOPROTEIN CHOLESTEROL, APOLIPOPROTEIN B, TRIGLYCERIDES, AND TO INCREASE HDL-C IN ADULT PATIENTS WITH PRIMARY HYPERLIPIDEMIA OR MIX DYSLIPIDEMIA

U-997 TREATMENT OF MAJOR DEPRESSIVE DISORDER BY DOSING AT INTERVALS OF 24 HOURS

U-998 ADJUNCTIVE THERAPY TO DIET TO REDUCE ELEVATED TOTAL CHOLESTEROL, LOW-DENSITY LIPOPROTEIN CHOLESTEROL, APOLIPOPROTEIN B, TRIGLYCERIDES AND TO INCREASE HDL-C IN ADULT PATIENTS WITH PRIMARY HYPERLIPIDEMIA OR MIXED DYSLIPIDEMIA

U-999 TREATMENT OF CHRONIC HEPATITIS B IN ADULT PATIENTS

U-1000 ADJUNCTIVE THERAPY TO DIET IN PATIENTS WITH HYPERLIPIDEMIAS

U-1001 METHOD FOR DELIVERING DRUG TO LUNG OF MAMMAL, COMPRISING ADMINISTERING DRUG PRODUCT BY INHALATION. TREATING A MAMMAL HAVING A CONDITION CAPABLE OF TREATMENT BY INHALATION, COMPRISING ADMINISTERING TO THE LUNG THE DRUG PRODUCT BY INHALATION

U-1002 METHOD OF TREATING INFLAMMATORY CONDITIONS

U-1003 A METHOD OF MYOCARDIAL PERFUSION IMAGING AND INCREASING CORONARY BLOOD FLOW

U-1004 TREATMENT OF PATIENTS WITH RELAPSED OR REFRACTORY PERIPHERAL T-CELL LYMPHOMA

U-1005 METHOD OF TREATING A STAPHYLOCOCCAL INFECTION

U-1006 NEW COMBINATION PRODUCT FOR THE EARLY TREATMENT OF RECURRENT HERPES LABIALIS (COLD SORES) TO REDUCE THE LIKELIHOOD OF ULCERATIVE COLD SORES AND TO SHORTEN THE LESION HEALING TIME IN ADULTS AND ADOLESCENTS (12 YEARS OF AGE AND OLDER)

U-1007 METHOD OF TREATING GOUT FLARES

U-1008 APPLICATION OF ANTISEPTIC WITH MOISTURIZERS FOR SURGICAL AND HEALTHCARE PERSONNEL SKIN DISINFECTION

U-1009 METHOD FOR ADMINISTRATION OF TESTOSTERONE

U-1010 TO REDUCE BLOOD PHENYLALANINE LEVELS IN PATIENTS WITH HYPERPHENYLALANINEMIA DUE TO TETRA HYDROBIOPTERIN RESPONSIVE PHENYLKETONURIA. KUVAN SHOULD BE TAKEN ORALLY WITH FOOD TO INCREASE ABSORPTION

U-1011 USE OF GRANISETRON TRANSDERMAL SYSTEM TO TREAT/PREVENT CHEMOTHERAPY INDUCED NAUSEA AND VOMITING

U-1012 METHOD FOR TREATING INSOMNIA WHILE REDUCING THE RISK OF AN ADVERSE DRUG INTERACTION

U-1013 METHOD OF USING RIBAVIRIN IN COMBINATION WITH PEGYLATED INTERFERON ALPHA-2B TO TREAT PATIENTS WITH CHRONIC HEPATITIS C

U-1014 METHOD OF USING RIBAVIRIN IN COMBINATION WITH INTERFERON ALPHA-2B (PEGYLATED AND NONPEGYLATED) TO TREAT PATIENTS WITH CHRONIC HEPATITIS C

U-1015 TREATMENT OF PATIENTS WITH RELAPSED OR REFRACTORY PERIPHERAL T-CELL LYMPHOMA

U-1016 IN COMBINATION WITH OTHER ANTIRETROVIRAL AGENTS FOR THE TREATMENT OF HIV-1 INFECTION IN TREATMENT-EXPERIENCED ADULT PATIENTS, WHO HAVE EVIDENCE OF VIRAL REPLICATION AND HIV-1 STRAINS RESISTANT TO AN NNRTI AND OTHER ANTIRETROVIRAL AGENTS

U-1017 A METHOD OF TREATING NASAL AND NON-NASAL SYMPTOMS OF SEASONAL ALLERGIC RHINITIS

U-1018 TREATMENT OF PULMONARY HYPERTENSION BY INHALATION

U-1019 TREATMENT OF PULMONARY HYPERTENSION

U-1020 METHOD OF USING COLCHICINE FOR THE PROPHYLAXIS OF GOUT FLARES

U-1021 SHORT-TERM TREATMENT (4-8 WEEKS) OF ACTIVE BENIGN GASTRIC ULCER

U-1022 FOR THE PREPARATION OF SKIN PRIOR TO SURGERY; HELPS REDUCE BACTERIA THAT CAN POTENTIALLY CAUSE SKIN INFECTION

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-1023 TREATMENT OF ATROPHIC VAGINITIS DUE TO MENOPAUSE

U-1024 REDUCTION OF ELEVATED INTRAOCULAR PRESSURE IN PATIENTS WITH GLAUCOMA OR OCULAR HYPERTENSION WHO REQUIRE ADJUNCTIVE OR REPLACEMENT THERAPY DUE TO INADEQUATELY CONTROLLED IOP

U-1025 TREATING FREQUENT HEARTBURN

U-1026 A METHOD OF TREATING HUMAN SUFFERING FROM OR SUSCEPTIBLE TO PSYCHOSIS.

U-1027 REDUCTION OF ELEVATED PLASMA STEROL AND/OR STANOL LEVELS IN A MAMMAL

U-1028 A METHOD OF DISTRIBUTING SODIUM OXYBATE UNDER CONTROL OF A CENTRAL PHARMACY

U-1029 METHOD FOR TREATING ACUTE ELEVATIONS OF BLOOD PRESSURE IN HUMAN SUBJECT IN NEED THEREOF

U-1030 IMPROVEMENT OF WALKING IN PATIENTS WITH MULTIPLE SCLEROSIS (MS)

U-1031 IMPROVE RESPIRATORY SYMPTOMS IN CYSTIC FIBROSIS IN PATIENTS WITH PSEUDOMONAS AERUGINOSA

U-1032 USE OF ROSUVASTATIN CALCIUM FOR THE PRIMARY PREVENTION OF CARDIOVASCULAR DISEASE IN INDIVIDUALS WITHOUT CLINICALLY EVIDENT CORONARY HEART DISEASE BUT WITH INCREASED RISK FACTORS

U-1033 TOPICAL TREATMENT OF ACNE VULGARIS

U-1034 TREATMENT OF ATTENTION DEFICIT HYPERACTIVITY DISORDER (ADHD) IN ADULTS

U-1035 NONSTEROIDAL ANTI-INFLAMMATORY DRUG INDICATED FOR RELIEF OF MILD TO MODERATE ACUTE PAIN

U-1036 METHOD OF TREATING TYPE 2 DIABETES MELLITUS BY ADMINISTERING A DIPEPTIDYL PEPTIDASE-4 INHIBITOR IN COMBINATION WITH INSULIN

U-1037 METHOD OF TREATING TYPE 2 DIABETES MELLITUS BY ADMINISTERING A DIPEPTIDYL PEPTIDASE-IV INHIBITOR IN COMBINATION WITH A PPAR-GAMMA AGONIST

U-1038 METHOD OF TREATING TYPE 2 DIABETES MELLITUS BY ADMINISTERING A DIPEPTIDYL PEPTIDASE-4 INHIBITOR IN COMBINATION WITH METFORMIN AND A PPAR-GAMMA AGONIST

U-1039 METHOD OF TREATING TYPE 2 DIABETES MELLITUS BY ADMINISTERING A DIPEPTIDYL PEPTIDASE-4 INHIBITOR IN COMBINATION WITH METFORMIN

U-1040 INHIBITION OF THROMBIN IN A PATIENT

U-1041 TREATMENT OF DISORDERS RESPONSIVE TO GROWTH HORMONE

U-1042 METHOD FOR STIMULATING CORONARY VASODILATION FOR PURPOSES OF IMAGING THE HEART

U-1043 MANAGEMENT OF MODERATE TO SEVERE PAIN

U-1044 TOPICAL TREATMENT OF SCALP PSORIASIS

U-1045 MAINTENANCE TREATMENT IN PATIENTS WITH LOCALLY ADVANCED OR METASTATIC NSCLC WHO HAVE NOT PROGRESSED ON 1ST-LINE TREATMENT WITH PLATINUM-BASED CHEMOTHERAPY

U-1046 MAINTENANCE TREATMENT OF PATIENTS WITH LOCALLY ADVANCED OR METASTATIC NSCLC WHOSE DISEASE HAS NOT PROGRESSED AFTER FOUR CYCLES PLATINUM-BASED CHEMOTHERAPY

U-1047 TREATMENT OF BIOPSY-CONFIRMED, PRIMARY SUPERFICIAL BASAL CELL CARCINOMA (SBCC)

U-1048 WORKS THROUGH THE INDUCTION OF INTERFERON AND OTHER CYTOKINES

U-1049 PROPHYLAXIS OF ORGAN REJECTION IN ADULT PATIENTS AT LOW-MODERATE IMMUNOLOGIC RISK RECEIVING A RENAL TRANSPLANT

U-1050 USE OF METAXALONE FOR TREATMENT OF MUSCULOSKELETAL CONDITIONS

U-1051 TREATMENT OF OROPHARYNGEAL CANDIDIASIS

U-1052 RELIEF OF SIGNS AND SYMPTOMS OF ARTHRITIS AND RISK-REDUCTION OF NSAID-ASSOCIATED GASTRIC ULCER

U-1053 RISK-REDUCTION OF NSAID-ASSOCIATED GASTRIC ULCER

U-1054 ONYCHOMYCOSIS OF THE TOENAIL CAUSED BY TRICOPHYTON RUBRUM OR TRICHOPHYTON MENTAGROPHYTES, ONCE DAILY USE FOR 12 CONSECUTIVE WEEKS

U-1055 AN ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS WHO ARE ALREADY TREATED WITH A THIAZOLIDINEDIONE (TZD) AND METFORMIN OR WHO HAVE INADEQUATE GLYCEMIC CONTROL ON A TZD OR METFORMIN ALONE

U-1056 TREATMENT OF PAIN USING A NASAL SPRAY OF KETOROLAC TROMETHAMINE

U-1057 TREATMENT OF INFLAMMATION AND PAIN USING A NASAL SPRAY OF KETOROLAC TROMETHAMINE

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-1058 USE OF THALIDOMIDE IN COMBINATION WITH DEXAMETHASONE FOR THE TREATMENT OF PATIENTS WITH NEWLY DIAGNOSED MULTIPLE MYELOMA

U-1059 ADJUNCTIVE THERAPY TO DIET TO PATIENTS WITH HYPERTRIGLYCERIDEMIA

U-1060 ADJUNCTIVE THERAPY TO DIET IN PATIENTS WITH ELEVATED CHOLESTEROL AND/OR LIPID LEVELS

U-1061 ADJUNCTIVE THERAPY TO DIET IN PATIENTS WITH MIXED DYSLIPIDEMIA

U-1062 ADMINISTRATION OF APPROVED PRODUCT FOR TREATMENT OF ALZHEIMER'S DISEASE

U-1063 TREATMENT OF ONLY INFLAMMATORY LESIONS (PAPULES AND PUSTULES) OF ROSACEA

U-1064 TREATMENT OF BIPOLAR DISORDER AND SCHIZOPHRENIA

U-1065 METHOD OF TREATING ANDROGEN RESPONSIVE OR MEDICATED CONDITION IN A MAMMAL BY ADMINISTERING A SAFE & EFFECTIVE AMOUNT OF DUTASTERIDE OR A PHARMACEUTICALLY ACCEPTABLE SOLVATE THEREOF.. CONDITIONS INCLUDE BENIGN PROSTATIC HYPERTROPHY

U-1066 METHOD OF TREATING AN ANDROGEN RESPONSE OR MEDIATED DISEASE IN A MAMMAL BY ADMINISTERING AN EFFECTIVE ANDROGEN RESPONSIVE OR MEDICATED DISEASE AMOUNT OF DUTASTERIDE..CONDITIONS INCLUDE BENIGN PROSTATIC HYPERPLASIA

U-1067 TREATMENT OF CANCER

U-1068 TREATMENT OF ASTHMA

U-1069 A METHOD OF TREATING A PATIENT WITH A PRESCRIPTION DRUG USING AN EXCLUSIVE COMPUTER DATABASE IN A COMPUTER SYSTEM FOR DISTRIBUTION

U-1070 A METHOD TO CONTROL ABUSE OF A SENSITIVE DRUG BY CONTROLLING WITH A COMPUTER PROCESSOR THE DISTRIBUTION OF THE SENSITIVE DRUG VIA AN EXCLUSIVITY CENTRAL PHARMACY THAT MAINTAINS A CENTRAL DATABASE

U-1071 METHOD OF TREATING BLADDER DYSFUNCTION WITH ONCE A DAY TROSPIMUM SALT FORMULATION

U-1072 THE MANAGEMENT OF MODERATE TO SEVERE CHRONIC PAIN IN PATIENTS REQUIRING A CONTINUOUS, AROUND-THE-CLOCK OPIOID ANALGESIC FOR AN EXTENDED PERIOD OF TIME

U-1073 USE FOR THE TREATMENT OF ASTHMA AND COPD

U-1074 USE OF EXENATIDE MAY RESULT IN REDUCTION IN BODY WEIGHT

U-1075 USE FOR THE TREATMENT OF ASTHMA

U-1076 REDUCE CHRONIC SEVERE DROOLING (I.E., SIALORRHEA) IN PATIENTS WITH NEUROLOGIC CONDITIONS ASSOCIATED WITH PROBLEM DROOLING

U-1077 PRETREATMENT OF PATIENTS WITH VITAMIN B12 AND FOLIC ACID PRIOR TO PEMETREXED DISODIUM ADMINISTRATION

U-1078 TREATMENT OF ACNE

U-1079 REVLIMID (LENALIDOMIDE) IS INDICATED FOR THE TREATMENT OF PATIENTS WITH TRANSFUSION-DEPENDENT ANEMIA IN MYELODYSPLASTIC SYNDROMES (MDS)

U-1080 METHOD TO TREAT PULMONARY HYPERTENSION BY ADMINISTERING AMBRISENTAN TO A PATIENT

U-1081 LUMIGAN IS A PROSTAGLANDIN ANALOG INDICATED FOR THE REDUCTION OF ELEVATED INTRAOCULAR PRESSURE IN PATIENTS WITH OPEN ANGLE GLAUCOMA OR OCULAR HYPERTENSION

U-1082 USE OF A COMBINATION OF TOBRAMYCIN AND DEXAMETHASONE TO TREAT OCULAR INFLAMMATION WHERE AN INFECTION OR RISK OF INFECTION EXISTS

U-1083 ACUTE TREATMENT OF MIGRAINE ATTACKS, WITH OR WITHOUT AURA, AND THE TREATMENT OF CLUSTER HEADACHE EPISODES

U-1084 RELIEF OF THE INFLAMMATORY AND PRURITIC MANIFESTATIONS OF CORTICOSTEROID RESPONSIVE DERMATOSES IN PATIENTS 12 YEARS OF AGE OR OLDER

U-1085 METHOD FOR TREATING IRRITABLE BOWEL SYNDROME AND METHOD FOR TREATING ABDOMINAL DISCOMFORT ASSOCIATED WITH IRRITABLE BOWEL SYNDROME

U-1086 TREATMENT OF AUTOIMMUNE DISEASE

U-1087 DETECTION OF NON-MUSCLE INVASIVE PAPILLARY CANCER OF THE BLADDER BY PHOTODYNAMIC CYSTOSCOPY

U-1088 RELIEF OF MUSCLE SPASM

U-1089 INHIBITION OF THROMBIN

U-1090 LO LOESTRIN FE IS INDICATED FOR THE PREVENTION OF PREGNANCY IN WOMEN WHO ELECT TO USE ORAL CONTRACEPTIVES AS A METHOD OF CONTRACEPTION

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-1091 ASSESSMENT OF BRONCHIAL HYPERRESPONSIVENESS IN PATIENTS 6 YEARS OF AGE OR OLDER WHO DO NOT HAVE CLINICALLY APPARENT ASTHMA

U-1092 TREATMENT OF BREAST CANCER

U-1093 TREATMENT OF PSEUDOBULBAR AFFECT

U-1094 MANAGEMENT OF CHRONIC MUSCULOSKELETAL PAIN

U-1095 METHOD OF TREATING OCULAR INFLAMMATION

U-1096 TREATMENT OF PATIENTS WITH METASTATIC BREAST CANCER

U-1097 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS WHEN TREATMENT WITH BOTH SAXAGLIPTIN AND METFORMIN IS APPROPRIATE

U-1098 METHOD OF TREATING HYPERPARATHYROIDISM; METHOD OF TREATING HYPERCALCEMIA

U-1099 TREATMENT OF PAIN, INCLUDING NEUROPATHIC PAIN ASSOCIATED WITH DIABETIC PERIPHERAL NEUROPATHY, POSTHERPETIC NEURALGIA, AND FIBROMYALGIA

U-1100 REDUCTION OF EXCESS ABDOMINAL FAT IN HIV-INFECTED PATIENTS WITH LIPODYSTROPHY

U-1101 METHOD OF TREATING EXCESSIVE DAYTIME SLEEPINESS IN PATIENTS WITH NARCOLEPSY

U-1102 METHOD OF TREATING CATAPLEXY IN PATIENTS WITH NARCOLEPSY

U-1103 TESTOSTERONE REPLACEMENT THERAPY IN MALES FOR CONDITIONS ASSOCIATED WITH A DEFICIENCY OR ABSENCE OF ENDOGENOUS TESTOSTERONE

U-1104 USE OF TRAMADOL FOR THE MANAGEMENT OF MODERATE TO MODERATELY SEVERE CHRONIC PAIN

U-1105 TOPICAL TREATMENT OF HEAD LICE INFESTATION IN PATIENTS FOUR (4) YEARS OF AGE AND OLDER

U-1106 TREATING HYPERTRIGLYCERIDEMIAS WITH REDUCTION OF FOOD EFFECT

U-1107 TREATING HYPERCHOLESTEROLEMIAS WITH REDUCTION OF FOOD EFFECT

U-1108 TREATING TYPE 2 DIABETES MELLITUS WITH EXENATIDE BY STIMULATING INSULIN RELEASE

U-1109 TREATMENT OF CUTANEOUS MANIFESTATIONS OF ERYTHEMA NODOSUM LEPROSUM (ENL) IN CONNECTION WITH A SPECIAL PROGRAM APPROVED BY FDA CALLED "SYSTEM FOR THALIDOMIDE EDUCATION AND PRESCRIBING SAFETY" (S.T.E.P.S.)

U-1110 METHOD OF TREATING A PATIENT WITH A PRESCRIPTION DRUG USING A COMPUTER DATABASE IN A COMPUTER SYSTEM FOR DISTRIBUTION

U-1111 NONSTEROIDAL ANTI-INFLAMMATORY DRUG INDICATED FOR RELIEF OF MILD TO MODERATE ACUTE PAIN

U-1112 METHOD OF MR IMAGING OF A MAMMAL

U-1113 TREATMENT AND PROPHYLAXIS OF INFLUENZA

U-1114 TREATMENT WITH GABAPENTIN, INCLUDING TREATMENT OF NEUROPATHIC PAIN, INCLUDING NEUROPATHIC PAIN ASSOCIATED WITH POSTHERPETIC NEURALGIA

U-1115 TREATMENT TO REDUCE THE RISK OF COPD EXACERBATIONS IN PATIENTS WITH SEVERE COPD ASSOCIATED WITH CHRONIC BRONCHITIS AND A HISTORY OF EXACERBATIONS

U-1116 METHOD OF ADMINISTERING COLCHICINE TO FAMILIAL MEDITERRANEAN FEVER PATIENTS

U-1117 TREATMENT OF BREAST CANCER

U-1118 USE FOR THE TREATMENT OF CHRONIC OBSTRUCTIVE PULMONARY DISEASE (COPD), INCLUDING CHRONIC BRONCHITIS AND EMPHYSEMA

U-1119 CONTRAST AGENT FOR MAGNETIC RESONANCE IMAGING

U-1120 TO REDUCE GASTROINTESTINAL SIDE EFFECTS ADMINISTER WITH A MEAL; AS STARTING DOSE ADMINISTER ONCE DAILY WITH EVENING MEAL

U-1121 METHOD OF TREATING TRAVELERS' DIARRHEA

U-1122 TREATMENT OF SECONDARILY INFECTED TRAUMATIC SKIN LESIONS DUE TO S. AUREUS AND S. PYOGENES

U-1123 TREATMENT OF ALCOHOL DEPENDENCE

U-1124 PREVENTION OF RELAPSE TO OPIOID DEPENDENCE, FOLLOWING OPIOID DETOXIFICATION

U-1125 METHOD FOR THE DETECTION OF NEUROENDOCRINE TUMORS

U-1126 USE IN COMBINATION WITH PREDNISONE FOR THE TREATMENT OF PATIENTS WITH METASTATIC CASTRATION-RESISTANT PROSTATE CANCER WHO HAVE RECEIVED PRIOR CHEMOTHERAPY

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

CONTAINING DOCETAXEL

- U-1127 TREATMENT OF PATENT DUCTUS ARTERIOSUS
- U-1128 TREATMENT OF CHRONIC HEPATITIS C (CHC) GENOTYPE 1 INFECTION IN COMBINATION WITH PEGINTERFERON ALFA AND RIBAVIRIN IN ADULT PATIENTS (>=18 YEARS OF AGE) WITH COMPENSATED LIVER DISEASE
- U-1129 TREATMENT OF HYPERCHOLESTEROLEMIA BY DOSING ONCE PER DAY IN THE EVENING OR AT NIGHT, WITH PRETREATMENT WITH A FLUSH INHIBITING AGENT SUCH AS ASPIRIN
- U-1130 SECONDARY PREVENTION OF CARDIOVASCULAR EVENTS BY DOSING ONCE PER DAY IN THE EVENING OR A NIGHT WITH PRETREATMENT WITH A FLUSH INHIBITING AGENT SUCH AS ASPIRIN
- U-1131 TREATMENT OF HYPERTRIGLYCERIDEMIA BY DOSING ONCE PER DAY IN THE EVENING OR AT NIGHT, WITH PRETREATMENT WITH A FLUSH INHIBITING AGENT SUCH AS ASPIRIN
- U-1132 TREATMENT OF HYPERCHOLESTEROLEMIA BY DOSING ONCE PER DAY IN THE EVENING OR AT NIGHT
- U-1133 SECONDARY PREVENTION OF CARDIOVASCULAR EVENTS BY DOSING ONCE PER DAY IN THE EVENING OR AT NIGHT
- U-1134 TREATMENT OF HYPERTRIGLYCERIDEMIA BY DOSING ONCE PER DAY IN THE EVENING OR AT NIGHT
- U-1135 TREATMENT OF HYPERCHOLESTEROLEMIA BY DOSING ONCE PER DAY IN THE EVENING OR AT NIGHT, THROUGH REDUCTION OF LDL-C, TC, TG, LP(A) AND INCREASE OF HDL-C
- U-1136 SECONDARY PREVENTION OF CARDIOVASCULAR EVENTS BY DOSING ONCE PER DAY IN THE EVENING OR AT NIGHT, THROUGH REDUCTION OF LDL-C, TC, TG, LP(A), AND INCREASE OF HDL-C
- U-1137 TREATMENT OF HYPERTRIGLYCERIDEMIA BY DOSING ONCE PER DAY IN THE EVENING OR AT NIGHT, THROUGH REDUCTION OF LDL-C, TC, TG, LP(A), AND INCREASE OF HDL-C
- U-1138 TREATMENT OF PRIMARY AND MIXED DYSLIPIDEMIA BY DOSING ONCE PER DAY IN THE EVENING OR AT NIGHT
- U-1139 REDUCTION IN RISK OF RECURRENT NONFATAL MYOCARDIAL INFARCTION BY DOSING ONCE PER DAY IN THE EVENING OR AT NIGHT
- U-1140 REDUCTION IN ELEVATED TC AND LDL-C BY DOSING ONCE PER DAY IN THE EVENING OR AT NIGHT
- U-1141 REDUCTION IN TG BY DOSING ONCE PER DAY IN THE EVENING OR AT NIGHT
- U-1142 TREATMENT OF PRIMARY AND MIXED DYSLIPIDEMIA BY DOSING ONCE PER DAY IN THE EVENING OR AT NIGHT, WITH PRETREATMENT WITH A FLUSH INHIBITING AGENT SUCH AS ASPIRIN
- U-1143 REDUCTION IN RISK OF RECURRENT NONFATAL MYOCARDIAL INFARCTION BY DOSING ONCE PER DAY IN THE EVENING OR A T NIGHT, WITH PRETREATMENT WITH A FLUSH INHIBITING AGENT SUCH AS ASPIRIN
- U-1144 REDUCTION IN ELEVATED TC AND LDL-C BY DOSING ONCE PER DAY IN THE EVENING OR AT NIGHT, WITH PRETREATMENT WITH A FLUSH INHIBITING AGENT SUCH AS ASPIRIN
- U-1145 REDUCTION IN TG BY DOSING ONCE PER DAY IN THE EVENING OR AT NIGHT, WITH PRETREATMENT WITH A FLUSH INHIBITING AGENT SUCH AS ASPIRIN
- U-1146 REDUCTION IN TG WITH REDUCED FLUSHING BY DOSING ONCE PER DAY IN THE EVENING OR AT NIGHT
- U-1147 TREATMENT OF PRIMARY AND MIXED DYSLIPIDEMIA BY DOSING ONCE PER DAY IN THE EVENING OR AT NIGHT, THROUGH REDUCTION OF LDL-C, TC, TG, LP(A), AND INCREASE OF HDL-C
- U-1148 REDUCTION IN RISK OF RECURRENT NONFATAL MYOCARDIAL INFARCTION BY DOSING ONCE PER DAY IN THE EVENING OR AT NIGHT, THROUGH REDUCTION OF LDL-C, TC, TG, LP(A), AND INCREASE OF HDL-C
- U-1149 TREATMENT OF HYPERTRIGLYCERIDEMIA BY DOSING ONCE PER DAY IN THE EVENING OR AT NIGHT, WITH PRETREATMENT WITH A FLUSH INHIBITING AGENT SUCH AS ASPIRIN
- U-1150 TREATMENT OF HYPERCHOLESTEROLEMIA BY DOSING ONCE PER DAY IN THE EVENING OR AT NIGHT, THROUGH REDUCTION IN TOTAL-C, LDL-C, TG, LP(A), AND INCREASE OF HDL-C
- U-1151 TREATMENT OF HYPERTRIGLYCERIDEMIA BY DOSING ONCE PER DAY IN THE EVENING OR AT NIGHT, THROUGH REDUCTION IN TOTAL-C, LDL-C, LP(A), AND INCREASE OF HDL-C
- U-1152 CYANOCOBALAMIN ADMINISTRATION THROUGH NASAL INFUSION

PATENT AND EXCLUSIVITY TERMS

ADB 110 of 225

PATENT USE

- U-1153 IN COMBINATION WITH OTHER ANTIRETROVIRAL AGENTS, IS INDICATED FOR THE TREATMENT OF HUMAN IMMUNODEFICIENCY VIRUS TYPE 1 (HIV-1) INFECTION IN ANTIRETROVIRAL TREATMENT-NAIVE ADULT PATIENTS, AS SET FORTH IN THE LABELING, INCLUDING I&U SECTION
- U-1154 TREATMENT OF PROTEIN KINASE RELATED DISORDERS, SUCH AS GASTROINTESTINAL STROMAL TUMORS, RENAL CELL CARCINOMA AND ADVANCED PANCREATIC NEUROENDOCRINE TUMORS, WITH SUNITINIB
- U-1155 USE OF THALIDOMIDE IN TREATMENT OF CUTANEOUS MANIFESTATIONS OF ERYTHEMA NODOSUM LEPROSUM (ENL)
- U-1156 TO REDUCE BLOOD PHENYLALANINE (PHE) LEVELS IN PATIENTS WITH HYPERPHENYLALANINEMIA (HPA)
- U-1157 RELIEF OF SYMPTOMS ASSOCIATED WITH RESPIRATORY ALLERGIES IN ADULTS AND CHILDREN 2 YEARS OF AGE AND OLDER AND FOR THE RELIEF OF SYMPTOMS ASSOCIATED WITH HIVES (URTICARIA) IN ADULTS AND CHILDREN 6 YEARS OF AGE AND OLDER
- U-1158 RELIEF OF SYMPTOMS ASSOCIATED WITH RESPIRATORY ALLERGIES AND FOR THE RELIEF OF SYMPTOMS ASSOCIATED WITH HIVES (URTICARIA) IN ADULTS AND CHILDREN 6 YEARS OF AGE AND OLDER
- U-1159 RELIEF OF SYMPTOMS ASSOCIATED WITH RESPIRATORY ALLERGIES, SWELLING OF THE NASAL PASSAGES AND SINUS CONGESTION AND PRESSURE IN ADULTS AND CHILDREN 12 YEARS OF AGE AND OLDER
- U-1160 RELIEF OF SYMPTOMS ASSOCIATED WITH RESPIRATORY ALLERGIES AND FOR THE RELIEF OF SYMPTOMS ASSOCIATED WITH HIVES (URTICARIA) IN ADULTS AND CHILDREN 6 YEARS OF AGE AND OLDER AND 12 YEARS OF AGE AND OLDER
- U-1161 FOR THE TREATMENT AND PROPHYLAXIS OF GOUT FLARES & THE TREATMENT OF FAMILIAL MEDITERRANEAN FEVER
- U-1162 TREATMENT OF SEBORRHEIC DERMATITIS OF THE SCALP
- U-1163 METHOD OF TREATING THROMBOSIS
- U-1164 METHOD OF TREATING AN ARGATROBAN TREATABLE CONDITION
- U-1165 USE FOR THE TREATMENT OF MULTIPLE MYELOMA
- U-1166 A METHOD FOR TREATMENT OF GOUT FLARES DURING PROPHYLAXIS
- U-1167 PROPHYLAXIS OF DEEP VEIN THROMBOSIS (DVT)
- U-1168 THE LONG TERM, ONCE-DAILY MAINTENANCE BRONCHODILATOR TREATMENT OF AIRFLOW OBSTRUCTION IN PATIENTS WITH CHRONIC OBSTRUCTIVE PULMONARY DISEASE (COPD), INCLUDING CHRONIC BRONCHITIS AND/OR EMPHYSEMA
- U-1169 MANAGEMENT OF BREAKTHROUGH PAIN IN CANCER PATIENTS 18 YEARS OF AGE AND OLDER WHO ARE RECEIVING AND TOLERANT TO OPIOID THERAPY FOR THEIR UNDERLYING PERSISTENT CANCER PAIN
- U-1170 TREATMENT OF HIV-1 INFECTION IN PEDIATRIC PATIENTS 12 YEARS OF AGE AND OLDER
- U-1171 REDUCTION OF THE RATE OF THROMBOTIC EVENTS IN PATIENTS WITH ACUTE CORONARY SYNDROME
- U-1172 TO REDUCE ELEVATED TOTAL-C, APO B, AND NON-HDL-C IN PATIENTS WITH PRIMARY HYPERLIPIDEMIA BY ADMINISTRATION OF EZETIMIBE IN COMBINATION WITH A STATIN
- U-1173 TO REDUCE ELEVATED TOTAL-C, LDL-C, APO B AND NON-HDL-C IN PATIENTS WITH PRIMARY HYPERLIPIDEMIA BY ADMINISTRATION OF EZETIMIBE ALONE OR IN COMBINATION WITH A STATIN OR WITH FENOFIBRATE
- U-1174 ADMINISTRATION OF REMODULIN DILUTED FOR INTRAVENOUS INFUSION WITH STERILE WATER FOR INJECTION, 0.9% SODIUM CHLORIDE INJECTION, OR FLOLAN STERILE DILUENT FOR INJECTION PRIOR TO ADMINISTRATION
- U-1175 REDUCTION OF CARDIAC TISSUE DAMAGE ASSOCIATED WITH MYOCARDIAL INFARCTION
- U-1176 TREATMENT OR PREVENTION OF STROKE
- U-1177 REDUCTION OF CARDIAC TISSUE DAMAGE ASSOCIATED WITH MYOCARDIAL INFARCTION
- U-1178 RELIEF OF MODERATE TO SEVERE CHRONIC PAIN
- U-1179 TREATMENT OF A CANCER MEDIATED BY AN ANAPLASTIC LYMPHOMA KINASE (ALK)
- U-1180 TREATMENT OF THE FOLLOWING INFECTIONS: COMPLICATED SKIN AND SKIN STRUCTURE INFECTIONS AND STAPHYLOCOCCUS AUREUS BLOODSTREAM INFECTIONS (BACTEREMIA) INCLUDING THOSE WITH RIGHT-SIDED INFECTIVE ENDOCARDITIS

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-1181 A METHOD OF TREATING OR PREVENTING OCULAR PAIN IN A PATIENT

U-1182 TREATMENT OF CYCLIC HEAVY MENSTRUAL BLEEDING

U-1183 A METHOD FOR ADMINISTERING FOLLICLE STIMULATING HORMONE (FSH) FOR OVARIAN FOLLICLE OR TESTICULAR STIMULATION IN THE HUMAN

U-1184 TREATMENT OF ERECTILE DYSFUNCTION AND THE SIGNS AND SYMPTOMS OF BENIGN PROSTATIC HYPERPLASIA

U-1185 TREATMENT OF OPIOID-INDUCED CONSTIPATION

U-1186 ADMINISTRATION OF AN INHALABLE POWDER COMPRISING TIOTROPIUM VIA DEVICE

U-1187 TREATMENT OF PATHOLOGICAL STATE BY ANTAGONIZING BRADYKININ RECEPTOR INCLUDING TREATMENT OF ACUTE ATTACKS OF HEREDITARY ANGIOEDEMA (HAE)

U-1188 METHOD OF TREATING TYPE 2 DIABETES MELLITUS IN PATIENTS FOR WHOM TREATMENT WITH BOTH SITAGLIPTIN AND SIMVASTATIN IS APPROPRIATE

U-1189 METHOD OF TREATING TYPE 2 DIABETES MELLITUS IN PATIENTS FOR WHOM TREATMENT WITH BOTH SITAGLIPTIN AND SIMVASTATIN IS APPROPRIATE, IN COMBINATION WITH METFORMIN

U-1190 METHOD OF TREATING TYPE 2 DIABETES MELLITUS IN PATIENTS FOR WHOM TREATMENT WITH BOTH SITAGLIPTIN AND SIMVASTATIN IS APPROPRIATE, IN COMBINATION WITH INSULIN

U-1191 METHOD OF TX TYPE 2 DM IN PTS FOR WHOM TREATMENT WITH BOTH SITAGLIPTIN AND SIMVASTATIN IS APPROPRIATE, IN COMBO WITH AN AGENT ACTING ON AN ATP-DEPENDENT CHANNEL IN BETA CELLS SUCH AS A SULFONYLUREA (INCL GLIPIZIDE, GLIMEPIRIDE & GLYBURIDE)

U-1192 METHOD OF TREATING TYPE 2 DIABETES MELLITUS IN PATIENTS FOR WHOM TREATMENT WITH BOTH SITAGLIPTIN AND SIMVASTATIN IS APPROPRIATE, IN COMBINATION WITH A SULFONYLUREA (SUCH AS GLIPIZIDE, GLIMEPIRIDE AND GLYBURIDE)

U-1193 METHOD OF TREATING TYPE 2 DIABETES MELLITUS IN PATIENTS FOR WHOM TREATMENT WITH BOTH SITAGLIPTIN AND SIMVASTATIN IS APPROPRIATE, IN COMBINATION WITH A PPAR-GAMMA AGONIST (SUCH AS PIOGLITAZONE AND ROSIGLITAZONE)

U-1194 METHOD FOR TREATING INSOMNIA

U-1195 PREVENTION AND TREATMENT OF SECONDARY HYPERPARATHYROIDISM ASSOCIATED WITH CHRONIC KIDNEY DISEASE (CKD) STAGE 5, WHICH MAY RESULT IN RENAL OSTEODYSTROPHY, WHILE AVOIDING HYPERPHOSPHATEMIA

U-1196 RELIEF OF SIGNS AND SYMPTOMS OF RHEUMATOID ARTHRITIS AND OSTEOARTHRITIS AND TO DECREASE RISK OF DEVELOPING UPPER GASTROINTESTINAL ULCERS IN PATIENTS WHO ARE TAKING IBUPROFEN FOR THOSE INDICATIONS

U-1197 METHOD OF TREATMENT OF CHILDREN WITH CENTRAL PRECOCIOUS PUBERTY

U-1198 RECTIV IS A NITRATE VASODILATOR INDICATED FOR THE TREATMENT OF MODERATE TO SEVERE PAIN ASSOCIATED WITH CHRONIC ANAL FISSURE

U-1199 TREATMENT AND PREVENTION OF POSTMENOPAUSAL OR GLUCOCORTICOID-INDUCED OSTEOPOROSIS AND TREATMENT TO INCREASE BONE MASS IN MEN WITH OSTEOPOROSIS

U-1200 REDUCING THE RISK OF STROKE AND SYSTEMIC EMBOLISM

U-1201 FOR THE TREATMENT OF INTERMEDIATE OR HIGH-RISK MYELOFIBROSIS

U-1202 METHOD FOR RELIEVING OR TREATING CONSTIPATION IN A PATIENT WITH IRRITABLE BOWEL SYNDROME

U-1203 METHOD FOR RELIEVING OR TREATING CONSTIPATION IN A HUMAN CONSTIPATED PATIENT

U-1204 TREATMENT OF UVEITIS

U-1205 TREATMENT OF MACULAR EDEMA

U-1206 DELIVERING AN OCULAR IMPLANT AS DESCRIBED IN THE DOSAGE AND ADMINISTRATION SECTION OF THE APPROVED LABELING OF OZURDEX

U-1207 INFANT USE AGED 1 MONTH TO LESS THAN ONE YEAR, GERD AND EROSIIVE ESOPHAGITIS

U-1208 TREATMENT OF HYPOTRICHOSIS OF THE EYELASHES BY INCREASING THEIR GROWTH INCLUDING LENGTH, THICKNESS AND DARKNESS

U-1209 TREATMENT OF HUMAN IMMUNODEFICIENCY VIRUS (HIV) INFECTION IN ADULT PATIENTS, AND TREATMENT OF HUMAN IMMUNODEFICIENCY VIRUS (HIV) INFECTION IN PEDIATRIC PATIENTS 3 YEARS OF AGE AND OLDER

U-1210 USE OF REVLIMID (LENALIDOMIDE) WHILE PREVENTING THE EXPOSURE OF A FETUS OR OTHER CONTRAINDICATED INDIVIDUAL TO REVLIMID (LENALIDOMIDE)

PATENT AND EXCLUSIVITY TERMS

ADB 112 of 225

PATENT USE

U-1211 USE OF REVLIMID (LENALIDOMIDE) TO INHIBIT THE SECRETION OF PRO-INFLAMMATORY CYTOKINES, INCLUDING TUMOR NECROSIS FACTOR ALPHA

U-1212 USE OF REVLIMID (LENALIDOMIDE) FOR THE TREATMENT OF MULTIPLE MYELOMA AND TRANSFUSION-DEPENDENT ANEMIA IN MYELOYDYSPLASTIC SYNDROMES (MDS)

U-1213 TOPICAL TREATMENT OF SEBORRHEIC DERMATITIS IN IMMUNOCOMPETENT PATIENTS 12 YEARS OF AGE AND OLDER

U-1214 METHOD FOR RELIEVING CONSTIPATION IN A HUMAN PATIENT THAT COMPRISES ADMINISTERING TO THE PATIENT A DOSAGE UNIT COMPRISING (I) 24MCG+/- 10% OF A DRUG SUBSTANCE AND (II) A PHARMACEUTICALLY SUITABLE EXCIPIENT

U-1215 USE OF REVLIMID (LENALIDOMIDE) FOR THE TREATMENT OF TRANSFUSION-DEPENDENT ANEMIA IN MYELOYDYSPLASTIC SYNDROMES (MDS)

U-1216 USE OF REVLIMID (LENALIDOMIDE) FOR THE TREATMENT OF MULTIPLE MYELOMA

U-1217 METHOD OF INCREASING HAIR GROWTH

U-1218 METHOD OF STIMULATING HAIR GROWTH

U-1219 METHOD OF INCREASING THE NUMBER OF HAIRS

U-1220 TREATMENT OF RENAL CELL CARCINOMA

U-1221 TO STIMULATE THE IMMUNE SYSTEM TO INDUCE T CELL PROLIFERATION

U-1222 TO INHIBIT THE PROLIFERATIVE ACTIVITY OF NEOPLASTIC CELLS

U-1223 METHOD FOR TREATING TYPE 2 DIABETES USING A SUSTAINED-RELEASE COMPOSITION CONTAINING EXENATIDE

U-1224 REDUCTIONS IN BODY WEIGHT ARE OBSERVED WITH EXENATIDE

U-1225 ACCELERATING THE TIME TO UPPER AND LOWER GASTROINTESTINAL RECOVERY FOLLOWING PARTIAL LARGE OR SMALL BOWEL RESECTION SURGERY WITH PRIMARY ANASTOMOSIS

U-1226 A METHOD OF PROVIDING A PREDETERMINED CONCENTRATION OF NITRIC OXIDE TO A PATIENT

U-1227 METHOD OF TREATING TYPE 2 DIABETES MELLITUS IN PATIENTS FOR WHOM TREATMENT WITH BOTH SITAGLIPTIN AND METFORMIN HCL EXTENDED RELEASE IS APPROPRIATE

U-1228 METHOD OF TREATING TYPE 2 DIABETES MELLITUS IN PATIENTS FOR WHOM TREATMENT WITH BOTH SITAGLIPTIN AND METFORMIN HCL EXTENDED RELEASE IS APPROPRIATE ALONE OR IN COMBINATION WITH INSULIN

U-1229 TREATMENT OF MILDLY TO MODERATELY ACTIVE ULCERATIVE COLITIS IN MALE PATIENTS

U-1230 A METHOD OF PROVIDING NITRIC OXIDE THERAPY TO A PATIENT

U-1231 TREATMENT OF MODERATE-TO-SEVERE PRIMARY RESTLESS LEG SYNDROME IN ADULTS

U-1232 USE AS ANTICOAGULANT IN PTS W/ UNSTABLE ANGINA UNDERGOING PTCA; W/ PROVISIONAL USE OF GLYCOPROTEIN IIB/IIIA INHIBITOR, AS ANTICOAGULANT IN PTS UNDERGOING PCI AND FOR PTS W/, OR AT RISK OF, HIT/HITTS UNDERGOING PCI.INTENDED FOR USE W/ASPIRIN

U-1233 TREATMENT OF CHRONIC HEPATITIS C (CHC) GENOTYPE 1 INFECTION, ADMINISTERED WITH FOOD

U-1234 FOR REDUCING TOTAL CHOLESTEROL (TOTAL-C), LDL-C, APO-LIPOPROTEIN B, OR TOTAL TRIGLYCERIDES, AND TREATING HYPERTRIGLYCERIDEMIA

U-1235 REDUCTION OF ELEVATED INTRAOCULAR PRESSURE IN PATIENTS WITH GLAUCOMA OR OCULAR HYPERTENSION

U-1236 USE OF THALOMID (THALIDOMIDE) FOR THE TREATMENT OF MULTIPLE MYELOMA

U-1237 COMBO W/ OTHER ANTIRETROVIRALS FOR TX OF HIV-1 IN ANTIRETROVIRAL TX-EXPERIENCED PT 6 YEARS UP, WHO HAVE EVIDENCE OF VIRAL REPLICATION AND HIV-1 STRAINS RESISTANT TO NON-NUCLEOSIDE REVERSE TRANSCRIPTASE INHIBITOR AND OTHER ANTIRETROVIRALS

U-1238 TREATMENT OF ANEMIA DUE TO CHRONIC KIDNEY DISEASE

U-1239 MAGNETIC RESONANCE IMAGING OF THE LIVER

U-1240 TREATMENT OF HEAVY MENSTRUAL BLEEDING IN WOMEN WITHOUT ORGANIC PATHOLOGY WHO CHOOSE TO USE AN ORAL CONTRACEPTIVE AS THEIR METHOD OF CONTRACEPTION

U-1241 MANAGEMENT OF MODERATE TO SEVERE PAIN BY ORALLY ADMINISTERING AN INTACT COMPOSITION AS CLAIMED

U-1242 PREVENTION OF RESPIRATORY DISTRESS (RDS) IN PREMATURE INFANTS

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

- U-1243 WITH DRY HANDS, GENTLY REMOVE THE SUPRENZA (PHENTERMINE HYDROCHLORIDE ODT) TABLET FROM THE BOTTLE. IMMEDIATELY PLACE THE SUPRENZA TABLET ON TOP OF THE TONGUE WHERE IT WILL DISSOLVE, THEN SWALLOW WITH OR WITHOUT WATER
- U-1244 METHOD OF TREATING TYPE 2 DIABETES MELLITUS BY ADMINISTERING A DIPEPTIDYL PEPTIDASE-4 INHIBITOR IN COMBINATION WITH SULFONYLUREA
- U-1245 METHOD OF TREATING TYPE 2 DIABETES MELLITUS BY ADMINISTERING A DIPEPTIDYL PEPTIDASE-4 INHIBITOR IN COMBINATION WITH PIOGLITAZONE
- U-1246 SINGLE DOSE ADMINISTRATION INTO THE SURGICAL SITE TO PRODUCE POSTSURGICAL ANALGESIA
- U-1247 MANAGEMENT OF POSTHERPETIC NEURALGIA (PHN) IN ADULTS
- U-1248 USE OF TOPICAL DICLOFENAC ON THE KNEE AND A SECOND TOPICAL MEDICATION ON THE SAME KNEE
- U-1249 TREATMENT OF MALE PATIENT HAVING A DISEASE OR CONDITION RESPONSIVE TO A TERATOGENIC DRUG
- U-1250 TREATMENT OF PAIN, INCLUDING NEUROPATHIC PAIN ASSOCIATED WITH DIABETIC PERIPHERAL NEUROPATHY OR SPINAL CORD INJURY, POSTHERPETIC NEURALGIA, AND FIBROMYALGIA
- U-1251 A METHOD OF CONTROLLING POSTOPERATIVE OCULAR PAIN AND BURNING/STINGING IN A PATIENT
- U-1252 METHOD FOR CHRONIC WEIGHT MANAGEMENT BY DECREASING FOOD INTAKE
- U-1253 METHOD FOR CHRONIC WEIGHT MANAGEMENT BY INDUCING SATIETY
- U-1254 METHOD FOR CHRONIC WEIGHT MANAGEMENT BY CONTROLLING WEIGHT GAIN
- U-1255 METHOD FOR CHRONIC WEIGHT MANAGEMENT BY TREATING OBESITY
- U-1256 TREATMENT OF SEBORRHEIC DERMATITIS
- U-1257 TREATMENT OF OPHTHALMIC DISORDERS
- U-1258 VISUALIZATION DURING VITRECTOMY PROCEDURES
- U-1259 PROPHYLAXIS OF HIV-1 INFECTION
- U-1260 TREATMENT OF PATIENTS WITH MULTIPLE MYELOMA WHO HAVE RECEIVED AT LEAST TWO PRIOR THERAPIES INCLUDING BORTEZOMIB AND AN IMMUNOMODULATORY AGENT AND HAVE DEMONSTRATED DISEASE PROGRESSION ON OR WITHIN 60 DAYS OF COMPLETION OF THE LAST THERAPY
- U-1261 REDUCTION OF THE RISK OF HOSPITALIZATION FOR ATRIAL FIBRILLATION
- U-1262 USE OF QSYMIA (PHENTERMINE AND TOPIRAMATE) FOR WEIGHT MANAGEMENT, INCLUDING, BUT NOT LIMITED TO EFFECTING WEIGHT LOSS, TREATING OBESITY, AND/OR TREATING OVERWEIGHT
- U-1263 TREATMENT OF CHRONIC OBSTRUCTIVE PULMONARY DISEASE (COPD) OR CHRONIC BRONCHITIS
- U-1264 TREATMENT OF A RESPIRATORY DISEASE
- U-1265 PATENTED METHOD OF USING REPAGLINIDE IN COMBINATION WITH METFORMIN AS INDICATED FOR IMPROVING GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS
- U-1266 METHOD OF TREATING MIDDLE-OF-THE-NIGHT INSOMNIA
- U-1267 TREATMENT OF RHEUMATOID ARTHRITIS BY DELAYED RELEASE FORMULATION OF 1MG OR 2MG OF PREDNISONE
- U-1268 TREATMENT OF PULMONARY, GASTROINTESTINAL AND/OR RHEUMATOLOGICAL DISEASES OR CONDITIONS BY USE OF DELAYED RELEASE FORMULATIONS OF 1MG OR 2MG PREDNISONE
- U-1269 TREATMENT OF RHEUMATOLOGIC, ALLERGIC, PULMONARY, GASTROINTESTINAL, DERMATOLOGIC DISEASES OR CONDITIONS BY THE USE OF A DELAYED RELEASE 5MG PREDNISONE TABLET
- U-1270 METHOD OF TREATING TYPE 2 DIABETES MELLITUS BY ADMINISTERING LINAGLIPTIN IN COMBINATION WITH INSULIN (WITH OR WITHOUT METFORMIN AND/OR PIOGLITAZONE)
- U-1271 TREATMENT OF ADULT PATIENTS WITH PHILADELPHIA CHROMOSOME-NEGATIVE (PH-) ACUTE LYMPHOBLASTIC LEUKEMIA (ALL) IN SECOND OR GREATER RELAPSE OR WHOSE DISEASE HAS PROGRESSED FOLLOWING TWO OR MORE ANTI-LEUKEMIA THERAPIES
- U-1272 TREATMENT OF SIGNS AND SYMPTOMS OF PARKINSON'S DISEASE BY APPLICATION OF CLAIMED TRANSDERMAL SYSTEM
- U-1273 TREATMENT OF RESTLESS LEGS SYNDROME BY APPLICATION OF CLAIMED TRANSDERMAL

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

DELIVERY SYSTEM

- U-1274 TREATMENT OF EXOCRINE PANCREATIC INSUFFICIENCY DUE TO CYSTIC FIBROSIS OR OTHER CONDITIONS
- U-1275 TREATMENT OF CHRONIC HEPATITIS B IN ADULTS AND PEDIATRIC PATIENTS 12 YEARS OF AGE AND OLDER
- U-1276 MANAGEMENT OF NEUROPATHIC PAIN ASSOCIATED WITH DIABETIC PERIPHERAL NEUROPATHY
- U-1277 METHOD OF INCREASING EYELASH GROWTH INCLUDING LENGTH, THICKNESS, DARKNESS AND/OR NUMBER OF EYELASHES BY ADMINISTERING BIMATOPROST TO AN EYELID MARGIN
- U-1278 METHOD OF TREATING IRRITABLE BOWEL SYNDROME WITH CONSTIPATION IN ADULTS
- U-1279 TREATMENT OF HIV INFECTION USING A COMPOSITION CONTAINING A PHARMACOKINETIC ENHANCER THAT INHIBITS CYTOCHROME P450 MONOOXYGENASE
- U-1280 USE OF A CALCIPOTRIENE CONTAINING FOAM FOR THE TREATMENT OF PSORIASIS
- U-1281 THE TREATMENT OF PATIENTS WITH METASTATIC CASTRATION-RESISTANT PROSTATE CANCER WHO HAVE PREVIOUSLY RECEIVED DOCETAXEL
- U-1282 PREVENTION OF ACUTE AND DELAYED NAUSEA AND VOMITING
- U-1283 A METHOD OF TREATING CHRONIC MYELOGENOUS LEUKEMIA
- U-1284 A METHOD OF TREATING A NEOPLASM
- U-1285 TREATMENT OF PATIENTS WITH RELAPSING FORMS OF MULTIPLE SCLEROSIS
- U-1286 A METHOD OF REDUCING THE RISK OF PULMONARY EDEMA IN PATIENTS IN NEED OF TREATMENT WITH INHALED NITRIC OXIDE
- U-1287 METHOD OF REDUCING TG LEVELS IN PATIENT SUFFERING FROM SEVERE HYPERTRIGLYCERIDEMIA
- U-1288 TREATMENT OF ERECTILE DYSFUNCTION BY ADMINISTERING A FILM-COATED TABLET
- U-1289 MANAGEMENT OF MODERATE TO SEVERE ACUTE PAIN
- U-1290 TREATMENT OF LUNG CANCER
- U-1291 TREATMENT OF ACUTE PROMYELOCYTIC LEUKEMIA (APL) IN PATIENTS WHOSE APL IS CHARACTERIZED BY THE PRESENCE OF THE (15;17) TRANSLOCATION OR PML/RAR-ALPHA GENE EXPRESSION
- U-1292 TREATMENT OF DISEASES OR CONDITIONS BY THE USE OF A DELAYED RELEASE 1, 2, OR 5 MG PREDNISONE TABLET
- U-1293 A METHOD OF LOWERING INTRAOCULAR PRESSURE IN A PATIENT WITH OPEN ANGLE GLAUCOMA OR OCULAR HYPERTENSION
- U-1294 METHOD OF TREATING GLAUCOMA IN A PATIENT
- U-1295 A METHOD OF TREATING A PATIENT WITH GLAUCOMA OR OCULAR HYPERTENSION
- U-1296 USE OF PEMETREXED WITH PRIOR AND/OR REPEATED VITAMIN B12 AND FOLIC ACID ADMINISTRATION
- U-1297 TREATMENT OF PULMONARY ARTERIAL HYPERTENSION BY INHIBITING ENDOTHELIN RECEPTORS
- U-1298 ADJUNCTIVE THERAPY IN THE TREATMENT OF PARTIAL SEIZURES
- U-1299 TREATMENT OF PATIENTS WITH LEUKEMIA INCLUDING CHRONIC MYELOID/MYELOGENOUS LEUKEMIA (CML)
- U-1300 TREATMENT OF PATIENTS WITH TYROSINE KINASE INHIBITOR (TKI) RESISTANT OR INTOLERANT CHRONIC MYELOID/MYELOGENOUS LEUKEMIA (CML)
- U-1301 TREATMENT OF DEEP VEIN THROMBOSIS (DVT)
- U-1302 TREATMENT OF PULMONARY EMBOLISM (PE)
- U-1303 REDUCTION IN THE RISK OF RECURRENCE OF DEEP VEIN THROMBOSIS (DVT) AND PULMONARY EMBOLISM
- U-1304 USE OF ONCE-A-DAY AMOXICILLIN PRODUCT TO TREAT TONSILLITIS AND/OR PHARYNGITIS SECONDARY TO STREPTOCOCCUS PYOGENES
- U-1305 TREATMENT OF HIV-1 INFECTION IN ADULT PATIENTS, AND TREATMENT OF HIV-1 INFECTION IN PEDIATRIC PATIENTS 3 YEARS OF AGE AND OLDER, CO-ADMINISTERED WITH RITONAVIR (PREZISTA/RITONAVIR) AND WITH OTHER ANTIRETROVIRAL AGENTS
- U-1306 TREATMENT OF THROMBOCYTOPENIA IN PATIENTS WITH CHRONIC HEPATITIS C TO ALLOW THE INITIATION AND MAINTENANCE OF INTERFERON-BASED THERAPY

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-1307 IN COMBINATION WITH OTHER ANTIRETROVIRAL AGENTS FOR THE TREATMENT OF HIV-1 INFECTION IN TREATMENT-NAIVE ADULT PATIENTS WITH HIV-1 RNA LESS THAN OR EQUAL TO 100,000 AT THE START OF THERAPY

U-1308 MULTIPLE MYELOMA

U-1309 BONE METASTASES

U-1310 FOR THE MAINTENANCE OF REMISSION OF ULCERATIVE COLITIS

U-1311 METHOD OF TREATING CYSTIC FIBROSIS

U-1312 USE FOR THE TREATMENT OF HYPERGLYCEMIA

U-1313 AS AN ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS

U-1314 USE IN COMBINATION WITH PREDNISONE FOR THE TREATMENT OF PATIENTS WITH METASTATIC CASTRATION-RESISTANT PROSTATE CANCER

U-1315 THE LONG TERM TREATMENT OF PROPHYLACTIC MANAGEMENT OF OCULAR HYPERTENSION AND GLAUCOMA

U-1316 A DOSING REGIMEN FOR THE TREATMENT OF HYPERCHOLESTEROLEMIA AND HYPERLIPIDEMIA IN PATIENTS WITH HOMOZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA USING AT LEAST THREE STEP-WISE INCREASING DOSES

U-1317 TREATMENT OF HYPERCHOLESTEROLEMIA, HYPERLIPIDEMIA AND HYPERLIPOPROTEINEMIA IN PATIENTS WITH HOMOZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA

U-1318 TREATMENT OF HYPERCHOLESTEROLEMIA BY DECREASING THE AMOUNT OR ACTIVITY OF MICROSOMAL TRIGLYCERIDE TRANSFER PROTEIN IN PATIENTS WITH HOMOZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA

U-1319 SYMPTOMATIC RELIEF OF NON-INFECTIOUS DIARRHEA

U-1320 TREATMENT OF ADULT PATIENTS WITH SHORT BOWEL SYNDROME WHO ARE DEPENDENT ON PARENTERAL SUPPORT

U-1321 TREATMENT OF PULMONARY MULTI-DRUG RESISTANT TUBERCULOSIS

U-1322 METHOD OF REDUCING OCULAR HYPERTENSION

U-1323 REDUCING THE RISK OF STROKE

U-1324 MANAGEMENT OF CYSTIC FIBROSIS PATIENTS

U-1325 INDUCTION OF REMISSION IN PATIENTS WITH ACTIVE, MILD TO MODERATE ULCERATIVE COLITIS

U-1326 METHOD OF INDUCING CONTRACEPTION IN A FEMALE OF REPRODUCTIVE AGE WHO HAS NOT YET REACHED PREMENOPAUSE

U-1327 METHOD FOR TREATING ACUTE MIGRAINE IN ADULTS, WITH OR WITHOUT AURA, COMPRISING IONTOPHORETIC TRANSDERMAL DELIVERY OF SUMATRIPTAN OR A SALT THEREOF, USING A FLOWABLE HYDROGEL FORMULATION

U-1328 METHOD FOR TREATING ACUTE MIGRAINE IN ADULTS, WITH OR WITHOUT AURA, COMPRISING IONTOPHORETIC TRANSDERMAL DELIVERY OF SUMATRIPTAN OR A SALT THEREOF

U-1329 TREATMENT OF PATIENTS WITH AN OVERACTIVE BLADDER

U-1330 METHODS OF TREATING LIPID METABOLISM AND GLYCOMETABOLISM DISORDERS COMPRISING ADMINISTERING AN INSULIN SENSITIVITY ENHANCER SUCH AS PIOGLITAZONE IN COMBINATION WITH AN INSULIN SECRETION ENHANCER

U-1331 METHODS OF REDUCING THE AMOUNT OF ACTIVE COMPONENTS ADMINISTERED TO A DIABETIC PATIENT COMPRISING ADMINISTERING AN INSULIN SENSITIVITY ENHANCER SUCH AS PIOGLITAZONE IN COMBINATION WITH AN INSULIN SECRETION ENHANCER

U-1332 METHODS OF REDUCING THE SIDE EFFECTS OF ACTIVE COMPONENTS ADMINISTERED TO A DIABETIC PATIENT COMPRISING ADMINISTERING AN INSULIN SENSITIVITY ENHANCER SUCH AS PIOGLITAZONE IN COMBINATION WITH AN INSULIN SECRETION ENHANCER

U-1333 METHODS OF LOWERING ELEVATED POST PRANDIAL BLOOD GLUCOSE LEVELS COMPRISING ADMINISTERING A DIPEPTIDYL PEPTIDASE INHIBITOR

U-1334 METHODS OF TREATING DIABETES COMPRISING ADMINISTERING AN INSULIN SENSITIVITY ENHANCER SUCH AS PIOGLITAZONE IN COMBINATION WITH AN INSULIN SECRETION ENHANCER

U-1335 METHODS OF MODIFYING GLUCOSE METABOLISM AND TREATING DIABETES COMPRISING ADMINISTERING A DIPEPTIDYL PEPTIDASE INHIBITOR AND ONE OR MORE OTHER THERAPEUTIC AGENTS SUCH AS METFORMIN

U-1336 METHODS OF TREATING DIABETES COMPRISING ADMINISTERING A DIPEPTIDYL PEPTIDASE

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

INHIBITOR AND METFORMIN

- U-1337 METHOD OF TREATING DIABETES COMPRISING ADMINISTERING ALOGLIPTIN
- U-1338 METHOD OF TREATING DIABETES COMPRISING ADMINISTERING A COMPOUND SUCH AS ALOGLIPTIN
- U-1339 METHODS OF TREATING DIABETES COMPRISING ADMINISTERING AN INSULIN SENSITIVITY ENHANCER SUCH AS PIOGLITAZONE IN COMBINATION WITH A BIGUANIDE SUCH AS METFORMIN
- U-1340 METHODS OF TREATING LIPID METABOLISM DISORDERS COMPRISING ADMINISTERING AN INSULIN SENSITIVITY ENHANCER SUCH AS PIOGLITAZONE IN COMBINATION WITH A BIGUANIDE SUCH AS METFORMIN
- U-1341 METHODS OF TREATING GLYCOMETABOLISM DISORDERS COMPRISING ADMINISTERING AN INSULIN SENSITIVITY ENHANCER SUCH AS PIOGLITAZONE IN COMBINATION WITH A BIGUANIDE SUCH AS METFORMIN
- U-1342 METHODS OF REDUCING THE AMOUNT OF ACTIVE COMPONENTS ADMINISTERED TO A DIABETIC PATIENT COMPRISING ADMINISTERING AN INSULIN SENSITIVITY ENHANCER SUCH AS PIOGLITAZONE IN COMBINATION WITH A BIGUANIDE SUCH AS METFORMIN
- U-1343 METHODS OF REDUCING THE SIDE EFFECTS OF ACTIVE COMPONENTS ADMINISTERED TO A DIABETIC PATIENT COMPRISING ADMINISTERING AN INSULIN SENSITIVITY ENHANCER SUCH AS PIOGLITAZONE IN COMBINATION WITH A BIGUANIDE SUCH AS METFORMIN
- U-1344 METHODS OF REDUCING THE SIDE EFFECTS OF ACTIVE COMPONENTS ADMINISTERED TO A DIABETIC PATIENT COMPRISING ADMINISTERING AN INSULIN SENSITIVITY ENHANCER SUCH AS PIOGLITAZONE IN COMBINATION WITH AN INSULIN PREPARATION
- U-1345 USE IN RELIEVING OR PREVENTING CONSTIPATION IN A HUMAN PATIENT WITH A DOSAGE UNIT COMPRISING 24MICROG+/- 10% OF A DRUG SUBSTANCE AND A PHARMACEUTICALLY SUITABLE EXCIPIENT
- U-1346 USE OF FEBUXOSTAT FOR THE MANAGEMENT OF HYPERURICEMIA IN PATIENTS SUFFERING FROM GOUT AND, WHEN USED WITH THEOPHYLLINE WITHOUT THE NEED FOR DOSE ADJUSTMENT OF THEOPHYLLINE
- U-1347 TREATMENT OF A SKIN DISORDER
- U-1348 TREATMENT OF OSTEOARTHRITIS
- U-1349 TREATMENT OF JUVENILE RHEUMATOID ARTHRITIS
- U-1350 TREATMENT OF ANKYLOSING SPONDYLITIS
- U-1351 TREATMENT OF ACUTE PAIN
- U-1352 TREATMENT OF PRIMARY DYSMENORRHEA
- U-1353 ADJUNCTIVE THERAPY TO LIPID-LOWERING MEDICATIONS AND DIET TO REDUCE LOW DENSITY LIPOPROTEIN-CHOLESTEROL, APOLIPOPROTEIN B, TOTAL CHOLESTEROL, AND NON-HIGH DENSITY LIPOPROTEIN CHOLESTEROL IN PTS WITH HOMOZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA
- U-1354 INHIBITION OF PREMATURE LH SURGES IN WOMEN UNDERGOING CONTROLLED OVARIAN HYPERSTIMULATION WITH FSH
- U-1355 MAINTENANCE TREATMENT OF ASTHMA AS PROPHYLACTIC THERAPY IN ADULT AND ADOLESCENT PATIENTS 12 YEARS OF AGE AND OLDER. PATENT CLAIMS METHOD FOR TREATING A RESPIRATORY DISEASE IN A CHILD
- U-1356 TREATMENT OF NASAL SYMPTOMS ASSOCIATED WITH SEASONAL ALLERGIC RHINITIS IN ADULTS AND CHILDREN 6 YEARS OF AGE AND OLDER. TREATMENT OF NASAL SYMPTOMS ASSOCIATED W PERENNIAL ALLERGIC RHINITIS IN ADULTS AND ADOLESCENTS 12 YEARS OF AGE AND OLDER
- U-1357 TREATMENT OF SYMPTOMS ASSOCIATED WITH SEASONAL AND PERENNIAL ALLERGIC RHINITIS IN ADULTS AND ADOLESCENTS 12 YEARS OF AGE AND OLDER. PATENT CLAIMS METHODS FOR TREATING A RESPIRATORY DISEASE IN A CHILD
- U-1358 TREATMENT OF BACTERIAL INFECTIONS IN THE NASAL PASSAGE OF ADULT PATIENTS AND HEALTH CARE WORKERS WITH METHICILLIN RESISTANT S. AUREUS
- U-1359 USE OF POMALIDOMIDE TO INHIBIT THE SECRETION OF PRO-INFLAMMATION CYTOKINES, INCLUDING TUMOR NECROSIS FACTOR ALPHA
- U-1360 USE OF POMALIDOMIDE FOR THE TREATMENT OF MULTIPLE MYELOMA
- U-1361 USE OF POMALIDOMIDE WHILE PREVENTING THE EXPOSURE OF A FETUS OR OTHER CONTRAINDICATED INDIVIDUAL TO POMALIDOMIDE
- U-1362 TREATMENT OF DISEASES OR CONDITIONS BY THE USE OF A DELAYED-RELEASE 1,2, OR 5MG PREDNISONE TABLET

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-1363 A METHOD OF TREATING OR PREVENTING OCULAR PAIN AND BURNING/STINGING FOLLOWING CORNEAL SURGERY

U-1364 MAINTENANCE TREATMENT OF MAJOR DEPRESSIVE DISORDER (MDD)

U-1365 PROPHYLAXIS OF ALLOGRAFT REJECTION IN ADULT PATIENTS RECEIVING A LIVER TRANSPLANT

U-1366 TREATMENT OF INFERTILITY THROUGH INDUCTION OF OVULATION AND PREGNANCY TO ANOVULATORY INFERTILE WOMEN

U-1367 METHOD OF ADMINISTERING FSH FOR THE TREATMENT OF INFERTILITY THROUGH INDUCTION OF OVULATION AND PREGNANCY IN ANOVULATORY INFERTILE WOMEN

U-1368 TREATMENT OF SOLID EXCRETORY SYSTEM TUMORS; ADVANCED RENAL CELL CARCINOMA (RCC), AFTER FAILURE OF TREATMENT WITH SUNITINIB OR SORAFENIB

U-1369 TREATMENT OF VAGINAL SYMPTOMS OF UROGENITAL ATROPHY BY ORALLY ADMINISTERING OSPEMIFENE WITH FOOD TO ENHANCE BIOAVAILABILITY OF OSPEMIFENE

U-1370 TREATMENT OF DYSpareunia ASSOCIATED WITH MENOPAUSE

U-1371 REDUCTION OF INTRAOCULAR PRESSURE IN PATIENTS WITH ELEVATED INTRAOCULAR PRESSURE OR GLAUCOMA

U-1372 ADMINISTRATION WITHOUT FOOD FOR TREATMENT OF HIV-1 INFECTION

U-1373 METHOD OF TREATING ACETAMINOPHEN OVERDOSE WITH ACETYLCYSTEINE SOLUTIONS

U-1374 TREATMENT OF PHILADELPHIA CHROMOSOME POSITIVE CHRONIC MYELOID LEUKEMIA (PH+CML)

U-1375 ADASUVE IS A TYPICAL ANTIPSYCHOTIC INDICATED FOR THE ACUTE TREATMENT OF AGITATION ASSOCIATED WITH SCHIZOPHRENIA OR BIPOLAR I DISORDER IN ADULTS

U-1376 TREATMENT OF INFLAMMATORY LESIONS OF NON-NODULAR MODERATE TO SEVERE ACNE VULGARIS

U-1377 IMPROVE RESPIRATORY SYMPTOMS IN CYSTIC FIBROSIS IN PATIENTS WITH PSEUDOMONAS AERUGINOSA

U-1378 TREATMENT OF A NITROGEN METABOLISM DISORDER

U-1379 IMPROVEMENT OF GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS WHO HAVE ONE OR MORE SPECIFIED CARDIOVASCULAR RISK FACTORS

U-1380 IMPROVEMENT OF GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS WHO HAVE ONE OR MORE SPECIFIED CARDIOVASCULAR RISK FACTORS WHEREIN THE PATIENT HAS CARDIOVASCULAR DISEASE

U-1381 USE OF PRASUGREL AND ASPIRIN IN PATIENTS REQUIRING THE REDUCTION OF THROMBOTIC CARDIOVASCULAR EVENTS

U-1382 TREATMENT OF NAUSEA AND VOMITING OF PREGNANCY IN WOMEN WHO DO NOT RESPOND TO CONSERVATIVE MANAGEMENT

U-1383 DOSAGE ADJUSTMENT OF A NITROGEN SCAVENGING DRUG IN THE TREATMENT OF A UREA CYCLE DISORDER

U-1384 METHOD OF TREATING MULTIPLE SCLEROSIS

U-1385 METHOD OF TREATING AN AUTOIMMUNE DISEASE SELECTED FROM AUTOIMMUNE POLYARTHRITIS AND MULTIPLE SCLEROSIS BUT NOT TREATING PSORIATIC ARTHRITIS

U-1386 A METHOD OF INCREASING THE TESTOSTERONE BLOOD LEVEL OF A PERSON IN NEED THEREOF

U-1387 REDUCTION IN RISK OF HOSPITALIZATION IN PATIENTS WITH A HISTORY OF PAROXYSMAL OR PERSISTENT AF WITHOUT SEVERE HEART FAILURE AND WITH ONE OR MORE RISK FACTORS BY ADMINISTRATION TWICE A DAY WITH MORNING AND EVENING MEALS

U-1388 TREATMENT OF PATIENTS WITH A HISTORY OF PAROXYSMAL OR PERSISTENT AF WITHOUT SEVERE HEART FAILURE AND WITH ONE OR MORE RISK FACTORS BY ADMINISTRATION TWICE A DAY WITH MORNING AND EVENING MEALS

U-1389 ELLA IS A PROGESTERONE AGONIST/ANTAGONIST EMERGENCY CONTRACEPTION INDICATED FOR THE PREVENTION OF PREGNANCY FOLLOWING UNPROTECTED INTERCOURSE OR A KNOWN OR SUSPECTED CONTRACEPTIVE FAILURE. ELLA CAN BE TAKEN WITH OR WITHOUT FOOD

U-1390 A METHOD OF INCREASING THE TESTOSTERONE BLOOD LEVEL OF AN ADULT MALE SUBJECT IN NEED THEREOF

U-1391 METHOD FOR TREATING OPIOID-INDUCED CONSTIPATION

U-1392 METHOD OF RELIEVING OR PREVENTING CONSTIPATION IN A HUMAN PATIENT WITH OPIOID-INDUCED CONSTIPATION

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

- U-1393 METHOD FOR RELIEVING OR TREATING CONSTIPATION IN A PATIENT WITH OPIOID-INDUCED CONSTIPATION
- U-1394 METHOD FOR RELIEVING CONSTIPATION IN A PATIENT WITH OPIOID-INDUCED CONSTIPATION THAT COMPRISES ADMINISTERING TO THE PATIENT A DOSAGE UNIT COMPRISING (I) 24MICROG +/- 10% OF A DRUG SUBSTANCE AND (II) A PHARMACEUTICALLY SUITABLE EXCIPIENT
- U-1395 USE IN RELIEVING OR PREVENTING CONSTIPATION IN A PATIENT WITH OPIOID-INDUCED CONSTIPATION WITH A DOSAGE UNIT COMPRISING 24MICROG +/- 10% OF A DRUG SUBSTANCE AND A PHARMACEUTICALLY SUITABLE EXCIPIENT
- U-1396 TREATMENT OF ADVANCED HORMONE RECEPTOR POSITIVE, HER2-NEGATIVE BREAST CANCER IN COMBINATION WITH EXEMESTANE AFTER FAILURE OF TREATMENT WITH LETROZOLE OR ANASTROZOLE
- U-1397 USE AS AN ANTISEPTIC FOR THE PREPARATION OF A PATIENT'S SKIN PRIOR TO SURGERY
- U-1398 METHOD OF TREATING CHRONIC HEPATITIS C
- U-1399 MANAGEMENT OF NEPHROPATHIC CYSTINOSIS BY ADMINISTERING A TOTAL DAILY DOSE IN TWO DIVIDED DOSES
- U-1400 FOR THE TREATMENT OF PRIMARY HYPERLIPIDEMIA, MIXED HYPERLIPIDEMIA OR HOMOZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA
- U-1401 INDICATED FOR LONG-TERM, ONCE-DAILY MAINTENANCE TREATMENT OF AIRFLOW OBSTRUCTION IN PTS WITH COPD, INCLUDING CHRONIC BRONCHITIS AND/OR EMPHYSEMA, ALSO TO REDUCE EXACERBATIONS OF COPD IN PTS WITH A HISTORY OF EXACERBATIONS
- U-1402 FOR USE IN THE TREATMENT OF PATIENTS WITH CHRONIC LYMPHOCYTIC LEUKEMIA (CLL) AND/OR INDOLENT B-CELL NON-HODGKIN LYMPHOMA (NHL)
- U-1403 FIRST-LINE TREATMENT OF METASTATIC NON SMALL-CELL LUNG CANCER (NSCLC) WITH EGFR EXON 19 DELETIONS OR EXON 21 (L858R) SUBSTITUTION MUTATIONS AS DETECTED BY AN FDA-APPROVED TEST
- U-1404 METHOD FOR TREATING CONSTIPATION IN A PATIENT WITH OPIOID-INDUCED CONSTIPATION BY OPENING CIC CHANNELS
- U-1405 THERAPEUTIC TREATMENT OF BONE METASTASES
- U-1406 TREATMENT OF MELANOMA
- U-1407 TREATMENT OF NEWLY DIAGNOSED PHILADELPHIA CHROMOSOME POSITIVE CHRONIC MYELOID LEUKEMIA (PH + CML)
- U-1408 TREATMENT OF PLAQUE PSORIASIS IN PATIENTS 18 YEARS OF AGE OR OLDER
- U-1409 TREATMENT OF HIV-1 BY ONCE DAILY ADMINISTRATION
- U-1410 TREATMENT OF CORTICOSTEROID-RESPONSIVE DERMATOSES
- U-1411 THIS DRUG IS ADMINISTERED BY SUBLINGUAL ROUTE TO HUMANS FOR MAINTENANCE TREATMENT OF OPIOID DEPENDENCE
- U-1412 TREATMENT OF ATOPIC DERMATITIS
- U-1413 ADMINISTRATION OF REMODULIN DILUTED FOR INTRAVENOUS INFUSION WITH FLOLAN STERILE DILUENT FOR INJECTION PRIOR TO INFUSION
- U-1414 USE OF REVLIMID (LENALIDOMIDE) FOR THE TREATMENT OF MANTLE CELL LYMPHOMA (MCL)
- U-1415 TREATING A PATIENT HAVING A CONDITION SUSCEPTIBLE TO TREATMENT WITH METHYLPHENIDATE, SUCH AS ADHD, BY ADMINISTERING THE FORMULATION RECITED IN CLAIMS 1 OR 2
- U-1416 USE OF FENOFIBRATE FOR REDUCING ELEVATED TOTAL CHOLESTEROL (TOTAL-C), LDL-C, APO-LIPOPROTEIN B, OR TOTAL TRIGLYCERIDES
- U-1417 USE FOR TREATMENT OF HELICOBACTER INFECTIONS
- U-1418 TREATMENT OF PATIENTS WITH UNRESECTABLE OR METASTATIC MELANOMA WITH BRAFV600E MUTATION AS DETECTED BY AN FDA APPROVED TEST
- U-1419 TREATMENT OF ACUTE CYANIDE POISONING THAT IS JUDGED TO BE LIFE THREATENING
- U-1420 METHOD OF ONCE A DAY ADMINISTRATION
- U-1421 SUBLINGUAL ADMINISTRATION OF A PHARMACEUTICAL COMPOSITION COMPRISING BUPRENORPHINE
- U-1422 METHOD OF TREATING PATIENTS NEEDING AN IRON SUPPLEMENT
- U-1423 AMYVID IS A RADIOACTIVE DIAGNOSTIC AGENT FOR POSITRON EMISSION TOMOGRAPHY (PET)

PATENT AND EXCLUSIVITY TERMS

ADB 119 of 225

PATENT USE

- IMAGING OF THE BRAIN TO ESTIMATE BETA-AMYLOID NEURITIC PLAQUE DENSITY IN ADULT PATIENTS WITH COGNITIVE IMPAIRMENT
- U-1424 LONG-TERM, ONCE DAILY MAINTENANCE TREATMENT OF AIRFLOW OBSTRUCTION IN PTS WITH COPD, INCLUDING CHRONIC BRONCHITIS AND/OR EMPHYSEMA, ALSO TO REDUCE EXACERBATIONS OF COPD IN PATIENTS WITH A HISTORY OF EXACERBATIONS
- U-1425 SUBLINGUAL ADMINISTRATION OF A PHARMACEUTICAL COMPOSITION COMPRISING BUPRENORPHINE AND NALOXONE
- U-1426 USE FOR TREATMENT OF DIAPER DERMATITIS COMPLICATED BY CANDIDIASIS
- U-1427 ALKYLATING DRUG INDICATED FOR THE TOPICAL TREATMENT OF STAGE IA AND IB MYCOSIS FUNGOIDES-TYPE CUTANEOUS T-CELL LYMPHOMA IN PATIENTS WHO HAVE RECEIVED PRIOR SKIN DIRECTED THERAPY
- U-1428 TOPICAL TREATMENT OF FACIAL ERYTHEMA OF ROSACEA
- U-1429 TREATMENT OF PATIENTS WITH BREAST CANCER WHOSE TUMORS OVEREXPRESS THE HER2 RECEPTOR
- U-1430 TREATMENT OF ALLERGIC RHINITIS, INCLUDING SEASONAL AND PERENNIAL ALLERGIC RHINITIS
- U-1431 METHOD OF TREATING HYPERGLYCEMIA TO IMPROVE GLYCEMIC CONTROL IN A PATIENT BY ORAL ADMIN OF ONCE A DAY OSMOTIC DOSAGE FORM OF GLIPIZIDE WITH POLYETHYLENE OXIDE, HYDROXYPROPYLMETHYLCELLULOSE, CELLULOSE ACETATE, AND SODIUM CHLORIDE
- U-1432 METHOD OF TREATMENT OF IRON-RELATED CONDITIONS WITH AT LEAST 0.6 GRAMS OF ELEMENTAL IRON VIA AN IRON CARBOHYDRATE COMPLEX
- U-1433 IMPROVEMENTS OF GLYCEMIC CONTROL IN INDIVIDUALS WITH TYPE 2 DIABETES WHO HAVE ONE OR MORE SPECIFIED CARDIOVASCULAR RISK FACTORS
- U-1434 TREATMENT OF PANCREATIC CANCER
- U-1435 COMBINATION USE OF TOPICAL DICLOFENAC ON THE KNEE AND ADMINISTRATION OF AN ORAL NSAID.
- U-1436 USE OF TOPICAL DICLOFENAC ON THE KNEE AND A SECOND TOPICAL AGENT SELECTED FROM SUNSCREEN AND INSECT REPELLANT
- U-1437 ADMINISTRATION OF REMODULIN DILUTED FOR INTRAVENOUS INFUSION WITH STERILE DILUENT FOR FLOLAN OR STERILE DILUENT FOR EPOPROSTENOL SODIUM PRIOR TO ADMINISTRATION
- U-1438 ZINGO INTRADERMAL INJECTION SYSTEM IS A DRUG DELIVERY SYSTEM THAT IS CAPABLE OF DELIVERING FINE DRY POWDERED LIDOCAINE HYDROCHLORIDE MONOHYDRATE FOR LOCAL ANESTHETIC ACTION
- U-1439 METHOD OF TREATING AN AFFECTIVE DISORDER SUCH AS DEPRESSION
- U-1440 USE OF INGENOL MEBUTATE TO TREAT ACTINIC KERATOSIS
- U-1441 A METHOD OF TREATING OR REDUCING OCULAR PAIN AND BURNING/STINGING
- U-1442 SUBCUTANEOUS INJECTION OF METHOTREXATE
- U-1443 ACCELERATING THE TIME TO UPPER AND LOWER GASTROINTESTINAL RECOVERY FOLLOWING SURGERIES THAT INCLUDE PARTIAL BOWEL RESECTION WITH PRIMARY ANASTOMOSIS
- U-1444 A DOSING REGIMEN OF AGGRASTAT (TIROFIBAN HYDROCHLORIDE) (25MCG/KG FOLLOWED BY 0.15MCG/KG/MIN INFUSION) TO REDUCE THE RATE OF THROMBOTIC CORONARY EVENTS ASSOCIATED WITH ACUTE CORONARY SYNDROME (ACS) IN PATIENTS WITH NON-ST ELEVATION ACS
- U-1445 METHOD OF TREATING PULMONARY ARTERIAL HYPERTENSION BY ADMINISTERING A PHARMACEUTICAL COMPOSITION COMPRISING MACITENTAN AND A POLYSORBATE, WHEREIN THE POLYSORBATE REPRESENTS 0.1 TO 1% OF THE WEIGHT OF SAID PHARMACEUTICAL COMPOSITION
- U-1446 METHOD OF TREATING PULMONARY HYPERTENSION COMPRISING ADMINISTERING MACITENTAN IN COMBINATION WITH A COMPOUND HAVING PHOSPHODIESTERASE-5 INHIBITORY PROPERTIES
- U-1447 TREATING PRIMARY HYPERCHOLESTEROLEMIA AND MIXED DYSLIPIDEMIA
- U-1448 TREATING SEVERE HYPERTRIGLYCERIDEMIA
- U-1449 METHOD OF ALLEVIATING A SKIN CONDITION
- U-1450 TREATMENT OF ALLERGIC RHINITIS SYMPTOMS
- U-1451 APPROVED INDICATIONS: APTIOM (ESLICARBAZEPINE ACETATE) IS INDICATED AS ADJUNCTIVE TREATMENT OF PARTIAL-ONSET SEIZURES AND APPROVED IN PATIENTS WITH

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

EPILEPSY. PATENT CLAIMS: IN A METHOD OF TREATING A SUBJECT AFFLICTED WITH EPILEPSY

U-1452 METHOD FOR CHRONIC WEIGHT MANAGEMENT

U-1453 A METHOD OF TREATING HYPOXIC RESPIRATORY FAILURE BY VERIFYING GAS INFORMATION OF NITRIC OXIDE PRIOR TO DELIVERY TO PATIENT

U-1454 PROPHYLAXIS OF INVASIVE ASPERGILLUS AND CANDIDA INFECTIONS

U-1455 TREATMENT OF PERIANAL WARTS

U-1456 TREATMENT OF MANTLE CELL LYMPHOMA

U-1457 A METHOD OF PURGING A NITRIC OXIDE DELIVERY SYSTEM

U-1458 A METHOD OF REDUCING INTRAOCULAR PRESSURE IN PATIENTS WITH OPEN ANGLE GLAUCOMA OR OCULAR HYPERTENSION

U-1459 TREATMENT OF CARCINOMA OF THE THYROID

U-1460 TREATMENT OF HERPES LABIALIS

U-1461 A METHOD OF GENERATING AN INJECTABLE FOAM OF CONTROLLED DENSITY AND BUBBLE SIZE

U-1462 A METHOD OF USING A SCLEROSING AGENT FOR THE TREATMENT OF INCOMPETENT GREAT SAPHENOUS VEINS, ACCESSORY SAPHENOUS VEINS AND VISIBLE VARICOSITIES OF THE GREAT SAPHENOUS (GSV) SYSTEM ABOVE AND BELOW THE KNEE

U-1463 A METHOD OF INTRAVENOUS INJECTION USING ULTRASOUND GUIDANCE, ADMINISTERED VIA A SINGLE CANNULA INTO THE LUMEN OF THE TARGET INCOMPETENT TRUNK VEINS OR BY DIRECT INJECTION INTO VARICOSITIES

U-1464 TREATMENT OF OPIOID DEPENDENCE/SUBLINGUAL OR BUCCAL APPLICATION

U-1465 USE OF THALIDOMIDE WHILE PREVENTING THE EXPOSURE OF A FETUS OR OTHER CONTRAINDICATED INDIVIDUAL TO THALIDOMIDE

U-1466 RELIEF OF SYMPTOMS ASSOCIATED WITH RESPIRATORY ALLERGIES ADULTS AND CHILDREN 6 YEARS OF AGE AND OLDER

U-1467 METHOD OF TREATING HEPATITIS C

U-1468 CONTROL OF PHOSPHOROUS LEVELS IN PATIENTS

U-1469 USE OF PHOSLYRA FOR REDUCTION OF SERUM PHOSPHOROUS IN PATIENTS

U-1470 FOR THE TREATMENT OF HEPATITIS C

U-1471 A METHOD FOR TREATING CARDIOVASCULAR DISEASE COMPRISING ADMINISTERING A RECONSTITUTED LYOPHILIZED PHARMACEUTICAL COMPOSITION COMPRISING EPOPROSTENOL, ARGININE AND SODIUM HYDROXIDE.

U-1472 INTENSIVE CARE UNIT SEDATION, INCLUDING SEDATION OF NON-INTUBATED PATIENTS PRIOR TO AND/OR DURING SURGICAL AND OTHER PROCEDURES

U-1473 MANAGEMENT OF RISK OF DRONEDARONE/BETA-BLOCKER INTERACTION IN PATIENTS IN SINUS RHYTHM WITH A HISTORY OF PAROXYSMAL OR PERSISTENT AF

U-1474 A METHOD FOR THE TREATMENT OF A PATIENT SUFFERING FROM A DISEASE TREATABLE WITH ROTIGOTINE, COMPRISING APPLYING THE CLAIMED TRANSDERMAL DELIVERY SYSTEM (TDS) TO THE SKIN OF THE PATIENT

U-1475 USE OF ORENITRAM FOR THE TREATMENT OF PULMONARY ARTERIAL HYPERTENSION (PAH) (WHO GROUP 1).

U-1476 INDICATED FOR THE LONG-TERM, ONCE-DAILY, MAINTENANCE TREATMENT OF AIRFLOW OBSTRUCTION IN PATIENTS WITH CHRONIC OBSTRUCTIVE PULMONARY DISEASE (COPD), INCLUDING CHRONIC BRONCHITIS AND/OR EMPHYSEMA.

U-1477 USE OF TOPICAL DICLOFENAC ON THE KNEE AND A SECOND TOPICAL PRESCRIPTION MEDICATION ON THE SAME KNEE

U-1478 METHOD OF REDUCING TG LEVELS IN PATIENT ON STATIN THERAPY SUFFERING FROM SEVERE HYPERTRIGLYCERIDEMIA

U-1479 INCREASE TEAR PRODUCTION TO TREAT PATIENTS WITH KERATOCONJUNCTIVITIS SICCA (DRY EYE).

U-1480 TREATMENT OF ADVANCED RENAL CELL CARCINOMA

U-1481 REDUCTION IN RISK OF OVERT HEPATIC ENCEPHALOPATHY (HE) RECURRENCE

U-1482 DICLOFENAC POTASSIUM FOR RELIEF OF MILD TO MODERATE ACUTE PAIN

U-1483 INCREASE TEAR PRODUCTION IN PATIENTS WITH KERATOCONJUNCTIVITIS SICCA (DRY EYE).

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-1484 COMBINATION PRODUCT FOR THE EARLY TREATMENT OF RECURRENT HERPES LABIALIS (COLD SORES) TO REDUCE THE LIKELIHOOD OF ULCERATIVE COLD SORES AND TO SHORTEN THE LESION HEALING TIME IN ADULTS AND CHILDREN (6 YEARS OF AGE AND OLDER)

U-1485 TREATING A SUBJECT UNDERGOING ABDOMINAL SURGERY BY ADMINISTERING ALVIMOPAN TO ACCELERATE THE TIME TO UPPER AND LOWER GASTROINTESTINAL RECOVERY FOLLOWING SURGERIES THAT INCLUDE PARTIAL BOWEL RESECTION WITH PRIMARY ANASTOMOSIS

U-1486 TREATMENT OF NON-24-HOUR SLEEP-WAKE DISORDER

U-1487 METHOD OF INCREASING EYELASH GROWTH

U-1488 USE OF TOPICAL DICLOFENAC FOR TREATING PAIN

U-1489 USE OF TOPICAL DICLOFENAC ON A JOINT FOR TREATING OSTEOARTHRITIS

U-1490 FOR USE IN PATIENTS HAVING SYMPTOMATIC OR PROGRESSIVE MEDULLARY THYROID CANCER, WITH UNRESECTABLE LOCALLY ADVANCED OR METASTATIC DISEASE

U-1491 TREATMENT OF CHRONIC LYMPHOCYTIC LEUKEMIA

U-1492 TREATMENT OF IRRITABILITY ASSOCIATED WITH AUTISTIC DISORDER

U-1493 METHOD FOR PREVENTING ITCHING ASSOCIATED WITH ALLERGIC CONJUNCTIVITIS

U-1494 SUBLINGUAL OR BUCCAL ADMINISTRATION OF A PHARMACEUTICAL COMPOSITION COMPRISING BUPRENORPHINE AND NALOXONE

U-1495 RISK REDUCTION OF REBLEEDING IN PTS FOLLOWING THERAPEUTIC ENDOSCOPY FOR ACUTE BLEEDING GASTRIC OR DUODENAL ULCERS IN ADULTS.

U-1496 METHOD TO TREAT HEMANGIOMA.

U-1497 NEURACEQ IS A RADIOACTIVE DIAGNOSTIC AGENT FOR POSITRON EMISSION TOMOGRAPHY (PET) IMAGING OF THE BRAIN TO ESTIMATE P-AMYLOID NEURITIC PLAQUE DENSITY IN ADULT PATIENTS WITH COGNITIVE IMPAIRMENT

U-1498 METHOD OF TREATING PATIENTS WITH GASTRIC RETENTIVE DOSAGE FORM

U-1499 MANAGEMENT OF ACUTE PAIN IN PATIENTS REQUIRING OPIOID ANALGESIA

U-1500 TESTOSTERONE REPLACEMENT THERAPY IN ADULT MALES FOR CONDITIONS ASSOCIATED WITH A DEFICIENCY OR ABSENCE OF ENDOGENOUS TESTOSTERONE; PRIMARY HYPOGONADISM (CONGENITAL OR ACQUIRED); HYPOGONADOTROPIC HYPOGONADISM (CONGENITAL OR ACQUIRED).

U-1501 PROPHYLAXIS OF DEEP VEIN THROMBOSIS AND PULMONARY EMBOLISM

U-1502 PROPHYLAXIS OF PULMONARY EMBOLISM

U-1503 METHOD OF TREATING TYPE 2 DIABETES MELLITUS BY ADMINISTERING LINAGLIPTIN IN COMBINATION WITH METFORMIN

U-1504 USE OF OTEZLA (APREMILAST) FOR INHIBITING PDE4

U-1505 USE OF OTEZLA (APREMILAST) FOR THE TREATMENT OF PSORIATIC ARTHRITIS

U-1506 TREATMENT OF PATIENTS WITH GASTROINTESTINAL STROMAL TUMOR (GIST), INCLUDING BUT NOT LIMITED TO PATIENTS PREVIOUSLY TREATED WITH IMATINIB AND PATIENTS WITH GIST HAVING RESISTANCE TO A KIT TYROSINE KINASE INHIBITOR

U-1507 TO MAINTAIN HEALING OF EE AND RELIEF OF HEARTBURN

U-1508 MANAGEMENT OF PAIN SEVERE ENOUGH TO REQUIRE DAILY, AROUND-THE-CLOCK, LONG TERM OPIOID TREATMENT BY ORALLY ADMINISTERING A PLURALITY OF COMPOSITE SUBUNITS AS CLAIMED

U-1509 TREATMENT OF FREQUENT HEARTBURN BY ADMINISTERING A GASTRIC ACID REDUCER

U-1510 MANAGEMENT OF PAIN SEVERE ENOUGH TO REQUIRE DAILY, AROUND-THE-CLOCK, LONG TERM OPIOID TREATMENT BY ORALLY ADMINISTERING AN INTACT COMPOSITION AS CLAIMED.

U-1511 TREATMENT OF HYPERTRIGLYCERIDEMIA

U-1512 REDUCTION OF THROMBOTIC CARDIOVASCULAR EVENTS

U-1513 TREATMENT OF HIV-1 INFECTION IN COMBINATION WITH OTHER ANTIRETROVIRAL AGENTS

U-1514 MANAGEMENT OF BREAKTHROUGH PAIN IN PATIENTS WITH CANCER BY BUCCAL OR SUBLINGUAL ADMINISTRATION OF FENTANYL

U-1515 METHOD OF TREATING IRRITABLE BOWEL SYNDROME WITH CONSTIPATION IN ADULT PATIENTS.

U-1516 METHOD OF TREATING CHRONIC IDIOPATHIC CONSTIPATION IN ADULT PATIENTS.

U-1517 TREATMENT OF BACTERIAL INFECTIONS USING A TWO-DOSE REGIMEN OF DALBAVANCIN.

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

- U-1518 MAINTAINING PUPIL SIZE BY PREVENTING INTRAOPERATIVE MIOSIS AND REDUCING POSTOPERATIVE OCULAR PAIN
- U-1519 METHOD FOR THE LONG TERM TREATMENT OF CHRONIC CONSTIPATION IN A HUMAN SUBJECT WITH IRRITABLE BOWEL SYNDROME
- U-1520 METHOD FOR THE LONG TERM TREATMENT OF CHRONIC CONSTIPATION IN A HUMAN SUBJECT
- U-1521 MAINTENANCE TREATMENT OF OPIOID DEPENDENCE
- U-1522 TREATMENT OF TYPE 2 DIABETES MELLITUS IN A PATIENT, WHEREIN GLYCEMIC CONTROL (HBA1C < 7.0%) IS NOT ACHIEVABLE USING ONE OR MORE OF INSULIN, METFORMIN, PIOGLITAZONE, OR ROSIGLITAZONE
- U-1523 METHOD OF INDUCING TOPICAL ANESTHESIA IN THE EYE
- U-1524 REDUCTION OF ELEVATED INTRAOCULAR PRESSURE
- U-1525 METHOD OF TREATING EXCESSIVE DAYTIME SLEEPINESS IN PATIENTS WITH NARCOLEPSY
- U-1526 THE TREATMENT OF PATIENTS WITH TRAVELERS' DIARRHEA (TD) OR THE REDUCTION IN RISK OF OVERT HEPATIC ENCEPHALOPATHY (HE) RECURRENCE
- U-1527 FOR THE TREATMENT OF OVERACTIVE BLADDER (OAB) WITH SYMPTOMS OF URGE URINARY INCONTINENCE, URGENCY, AND URINARY FREQUENCY
- U-1528 A METHOD OF LOWERING INTRAOCULAR PRESSURE
- U-1529 ADJUNCTIVE TREATMENT OF MAJOR DEPRESSIVE DISORDER (MDD)
- U-1530 USE OF ARIPIPRAZOLE IN EXTENDED RELEASE INJECTABLE SUSPENSION
- U-1531 METHOD FOR TRANSDERMAL DELIVERY OF TESTOSTERONE
- U-1532 METHOD OF TREATING EXCESSIVE DAYTIME SLEEPINESS AND/OR CATAPLEXY IN NARCOLEPSY PATIENTS WITH SODIUM OXYBATE WHEN DIVALPROEX SODIUM IS CONCOMITANTLY ADMINISTERED.
- U-1533 PULMONARY ADMINISTRATION OF PARTICLES COMPRISING A DIKETOPIPERAZINE AND INSULIN.
- U-1534 ADMINISTRATION OF A COMPOSITION COMPRISING INSULIN COMPLEXED WITH A DIKETOPIPERAZINE.
- U-1535 ADMINISTRATION OF A COMPOSITION COMPRISING INSULIN COMPLEXED WITH MICROPARTICLES OF A DIKETOPIPERAZINE.
- U-1536 ADMINISTRATION OF A COMPOSITION COMPRISING A DIKETOPIPERAZINE AND INSULIN.
- U-1537 TREATMENT OF A PATIENT HAVING DIABETES MELLITUS WITH A PRANDIAL RAPID ACTING INSULIN.
- U-1538 ADMINISTRATION OF FDKP MICROPARTICLES COMPRISING INSULIN.
- U-1539 PULMONARY ADMINISTRATION OF AN INSULIN COMPOSITION COMPRISING FDKP AT THE BEGINNING OF A MEAL TO A PATIENT ALSO BEING TREATED WITH A LONG-ACTING INSULIN.
- U-1540 BUTRANS IS A PARTIAL OPIOID AGONIST PRODUCT INDICATED FOR THE MANAGEMENT OF PAIN SEVERE ENOUGH TO REQUIRE DAILY, AROUND-THE-CLOCK, LONG TERM OPIOID TREATMENT FOR WHICH ALTERNATIVE TREATMENT OPTIONS ARE INADEQUATE.
- U-1541 TREATMENT OF PATIENTS WITH TUBEROUS SCLEROSIS COMPLEX (TSC) WHO HAVE SUBEPENDYMAL GIANT CELL ASTROCYTOMA (SEGA) THAT REQUIRES THERAPEUTIC INTERVENTION BUT CANNOT BE CURATIVELY RESECTED.
- U-1542 FOR USE IN THE TREATMENT OF PATIENTS WITH CHRONIC LYMPHOCYTIC LEUKEMIA AND/OR NON-HODGKINS LYMPHOMA
- U-1543 TREATMENT OF A PATIENT BY ADMINISTERING THE FORMULATION RECITED IN CLAIM 1 OR CLAIM 23
- U-1544 TREATMENT OF PATIENTS WITH RELAPSED OR REFRACTORY PERIPHERAL T-CELL LYMPHOMA (PTCL).
- U-1545 A METHOD OF TRANSDERMALLY DELIVERING TESTOSTERONE
- U-1546 FOR USE IN THE TREATMENT OF MALIGNANT HYPERTHERMIA IN CONJUNCTION WITH APPROPRIATE SUPPORTIVE MEASURES AND FOR THE PREVENTION OF MALIGNANT HYPERTHERMIA IN PATIENTS AT HIGH RISK.
- U-1547 TREATMENT OF CHRONIC OBSTRUCTIVE PULMONARY DISEASE (COPD), CHRONIC BRONCHITIS OR EMPHYSEMA
- U-1548 FOR THE LONG-TERM, ONCE-DAILY MAINTENANCE TREATMENT OF AIRFLOW OBSTRUCTION IN PATIENTS WITH COPD, INCLUDING CHRONIC BRONCHITIS AND/OR EMPHYSEMA, ALSO TO REDUCE EXACERBATIONS OF COPD IN PATIENTS WITH A HISTORY OF EXACERBATIONS

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-1549 FOR THE TREATMENT OF PATIENTS WITH RELAPSED CHRONIC LYMPHOCYTIC LEUKEMIA

U-1550 METHOD OF TREATING METASTATIC PAPILLARY RENAL CELL CARCINOMA WITH TEMSIROLIMUS.

U-1551 METHOD OF TREATING PAPILLARY RENAL CELL CARCINOMA WITH TEMSIROLIMUS, IN THE ABSENCE OF INTERFERON ALPHA.

U-1552 FOR HEALING OF ALL GRADES OF EROSIIVE ESOPHAGITIS (EE)

U-1553 TO MAINTAIN HEALING OF EE AND RELIEF OF HEARTBURN

U-1554 FOR THE TREATMENT OF HEARTBURN ASSOCIATED WITH SYMPTOMATIC NON-EROSIVE GASTROESOPHAGEAL DISEASE (GERD)

U-1555 MANAGEMENT OF MODERATE TO SEVERE PAIN SEVERE ENOUGH TO REQUIRE DAILY, AROUND-THE-CLOCK, LONG-TERM OPIOID TREATMENT AND FOR WHICH ALTERNATIVE TREATMENT OPTIONS ARE INADEQUATE.

U-1556 MANAGEMENT OF PAIN SEVERE ENOUGH TO REQUIRE DAILY, AROUND-THE-CLOCK, LONG-TERM OPIOID TREATMENT AND FOR WHICH ALTERNATIVE TREATMENT OPTIONS ARE INADEQUATE

U-1557 A METHOD OF TESTOSTERONE REPLACEMENT THERAPY COMPRISING THE STEP OF NASALLY ADMINISTERING TO A PATIENT IN NEED OF SUCH TREATMENT AN EFFECTIVE AMOUNT OF TESTOSTERONE GEL FORMULATION.

U-1558 FOR THE TREATMENT OF PATIENTS WITH RELAPSED FOLLICULAR B-CELL NON-HODGKIN LYMPHOMA OR [RELAPSED] SMALL LYMPHOCYTIC LYMPHOMA

U-1559 INDICATED FOR THE ONCE-DAILY MAINTENANCE TREATMENT OF ASTHMA AS PROPHYLACTIC THERAPY IN PATIENTS AGED 12 YEARS OF AGE AND OLDER

U-1560 A METHOD OF DISRUPTING LEUKOCYTE FUNCTION, INCLUDING AS AN INHIBITOR OF PI3KDELTA KINASE

U-1561 USE OF OTEZLA (APREMILAST) FOR THE TREATMENT OF PSORIATIC ARTHRITIS

U-1562 TREATMENT OF PATIENTS WITH HEPATIC ENCEPHALOPATHY (HE)

U-1563 A METHOD OF TRANSDERMAL ADMINISTRATION OF A PHYSIOLOGICALLY ACTIVE AGENT TO A SUBJECT.

U-1564 A METHOD OF TREATING GAUCHER'S DISEASE

U-1565 METHOD OF TREATING, AS INITIAL LOADING DOSE FOR MONOTHERAPY OR ADJUNCTIVE THERAPY, PARTIAL ONSET-SEIZURES IN A PATIENT WITH EPILEPSY AGED 17 YEARS OR OLDER WHEN ORAL ADMINISTRATION IS TEMPORARILY NOT FEASIBLE

U-1566 METHOD OF TREATING, AS MONOTHERAPY OR ADJUNCTIVE THERAPY, PARTIAL-ONSET SEIZURES IN A PATIENT WITH EPILEPSY AGED 17 YEARS AND OLDER

U-1567 METHOD OF TREATING, AS INITIAL LOADING DOSE FOR MONOTHERAPY OR ADJUNCTIVE THERAPY, PARTIAL ONSET-SEIZURES IN A PATIENT WITH EPILEPSY AGED 17 YEARS OR OLDER

U-1568 METHOD OF TREATING, AS MONOTHERAPY OR ADJUNCTIVE THERAPY, PARTIAL-ONSET SEIZURES IN A PATIENT WITH EPILEPSY AGED 17 YEARS AND OLDER WHEN ORAL ADMINISTRATION IS TEMPORARILY NOT FEASIBLE

U-1569 TREATMENT OF BACTERIAL SKIN AND SKIN STRUCTURE INFECTIONS

U-1570 TREATMENT OF BACTERIAL SKIN AND SKIN STRUCTURE INFECTIONS USING A SINGLE DOSE

U-1571 TREATMENT OF GAUCHER DISEASE TYPE 1

U-1572 TREATMENT OF HUMAN IMMUNODEFICIENCY VIRUS (HIV) INFECTION.

U-1573 USE OF RUXOLITINIB (JAKAFI) FOR INHIBITING JANUS ASSOCIATED KINASES (JAKS) JAK1 AND/OR JAK2.

U-1574 A METHOD OF CATALYZING THE HYDROLYSIS OF GLUCOCEREBROSIDE TO GLUCOSE AND CERAMIDE.

U-1575 PATIENTS WITH SEVERE APLASTIC ANEMIA WHO HAVE HAD AN INSUFFICIENT RESPONSE TO IMMUNOSUPPRESSIVE THERAPY

U-1576 TREATMENT OF LEUKEMIA

U-1577 CONTROL OF SERUM PHOSPHOROUS LEVELS

U-1578 TREATMENT OF ACUTE OTITIS MEDIA

U-1579 USE IN COMBINATION WITH PREDNISONE FOR THE TREATMENT OF PATIENTS WITH METASTATIC CASTRATION-RESISTANT PROSTATE CANCER

U-1580 USE IN COMBINATION WITH PREDNISONE FOR THE TREATMENT OF PATIENTS WITH METASTATIC CASTRATION-RESISTANT PROSTATE CANCER WHO HAD RECEIVED PRIOR DOCETAXEL

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

CHEMOTHERAPY

- U-1581 IN COMBINATION WITH DABRAFENIB FOR THE TREATMENT OF UNRESECTABLE OR METASTATIC MELANOMA.
- U-1582 TREATMENT OF UNRESECTABLE OR METASTATIC MELANOMA
- U-1583 FOR CHRONIC WEIGHT MANAGEMENT FOR TREATING OVERWEIGHT OR OBESITY
- U-1584 USE OF NALTREXONE AND BUPROPION IN A LAYERED FORMULATION FOR CHRONIC WEIGHT MANAGEMENT FOR AFFECTING WEIGHT LOSS
- U-1585 USE OF NALTREXONE AND BUPROPION BASED ON AN ESCALATING DOSE SCHEDULE
- U-1586 FOR EFFECT ON BLOOD GLUCOSE PARAMETERS IN PATIENTS WITH INSULIN RESISTANCE
- U-1587 SINGLE-DOSE INFILTRATION INTO THE SURGICAL SITE TO PRODUCE POSTSURGICAL ANALGESIA.
- U-1588 THE TREATMENT OF PATIENTS WITH METASTATIC CASTRATION-RESISTANT PROSTATE CANCER (CRPC).
- U-1589 METHOD OF USE FOR REDUCING BLOOD PHENYLALANINE LEVELS IN A HUMAN SUFFERING FROM HYPERPHENYLALANINEMIA
- U-1590 KUVAN IS INDICATED TO REDUCE BLOOD PHENYLALANINE LEVELS IN PATIENTS WITH HYPERPHENYLALANINEMIA
- U-1591 TREATMENT OF ASTHMA IN PATIENTS AGED 12 YEARS AND OLDER
- U-1592 TO REDUCE SERUM PHOSPHATE IN PATIENTS WITH END STAGE RENAL DISEASE
- U-1593 MAINTENANCE TREATMENT OF BRONCHOSPASM ASSOCIATED WITH COPD, INCLUDING CHRONIC BRONCHITIS AND EMPHYSEMA, AND REDUCTION OF EXACERBATIONS IN COPD PATIENTS.
- U-1594 DILATION OF THE PUPIL
- U-1595 USE OF OTEZLA (APREMILAST) FOR THE TREATMENT OF PSORIASIS
- U-1596 LAMICTAL IS AN ANTIEPILEPTIC DRUG (AED) INDICATED FOR: EPILEPSY-ADJUNCTIVE THERAPY IN PATIENTS GREATER THAN OR EQUAL TO 2 YEARS OF AGE: (1.1) PARTIAL SEIZURES PRIMARY GENERALIZED TONIC-CLONIC SEIZURES
- U-1597 TREATMENT OF DIABETIC MACULAR EDEMA
- U-1598 METHOD OF ADMINISTRATION OF CONTROLLED RELEASE OXYMORPHONE
- U-1599 MANAGEMENT OF MILD TO MODERATE PAIN, MANAGEMENT OF MODERATE TO SEVERE PAIN AS AN ADJUNCT TO OPIOID ANALGESICS, REDUCTION IN FEVER THROUGH ANTI-INFLAMMATORY, ANALGESIC, AND ANTIPYRETIC ACTIVITY
- U-1600 DOSAGE MODIFICATION FOLLOWING ELEVATED LIVER ENZYMES IN TREATMENT OF IDIOPATHIC PULMONARY FIBROSIS
- U-1601 DOSE ESCALATION OVER 14 DAYS FOR TREATMENT OF IDIOPATHIC PULMONARY FIBROSIS
- U-1602 METHOD OF ADMINISTERING PIRFENIDONE CAPSULES TO TREAT A FIBROTIC CONDITION
- U-1603 METHOD FOR ADMINISTERING PIRFENIDONE TO REDUCE DRUG INTERACTIONS WITH FLUVOXAMINE
- U-1604 METHOD FOR ADMINISTERING PIRFENIDONE TO REDUCE DRUG INTERACTIONS WITH A STRONG INHIBITOR OF CYP1A2
- U-1605 METHOD FOR ADMINISTERING PIRFENIDONE TO AVOID REDUCED EFFICACY BY DISCONTINUING SMOKING OR BY DISCONTINUING OR AVOIDING ANOTHER STRONG CYP1A2 INDUCER
- U-1606 METHOD FOR ADMINISTERING PIRFENIDONE WHILE AVOIDING OR DISCONTINUING CONCOMITANT USE OF A MODERATE TO STRONG INHIBITOR OF BOTH CYP1A2 AND ANOTHER CYP ENZYME INVOLVED IN PIRFENIDONE METABOLISM
- U-1607 METHOD OF ADMINISTERING A DOSAGE FORM THAT INCLUDES A GRANULATE FORMULATION OF PIRFENIDONE TO TREAT A FIBROTIC CONDITION
- U-1608 DOSE ESCALATION OVER 14 DAYS FOR TREATMENT OF A FIBROSIS CONDITION
- U-1609 CONTINUED DOSING OR DOSAGE MODIFICATION FOLLOWING ELEVATED LIVER ENZYMES IN TREATMENT OF IDIOPATHIC PULMONARY FIBROSIS
- U-1610 CONTINUED DOSING OR DOSAGE MODIFICATION FOLLOWING ELEVATED LIVER ENZYMES IN USE OF PIRFENIDONE
- U-1611 METHOD FOR ADMINISTERING PIRFENIDONE TO AVOID REDUCED EFFICACY BY DISCONTINUING SMOKING OR BY DISCONTINUING A STRONG CYP1A2 INDUCER
- U-1612 METHOD FOR ADMINISTERING PIRFENIDONE TO AVOID REDUCED EFFICACY BY AVOIDING

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

SMOKING OR BY AVOIDING ANOTHER STRONG CYP1A2 INDUCER

U-1613 DOSAGE MODIFICATION IN TREATMENT WITH PIRFENIDONE TO REDUCE DRUG INTERACTIONS WITH CIPROFLOXACIN

U-1614 USE OF TOPICAL DICLOFENAC SODIUM FOR TREATING PAIN

U-1615 FOR THE TREATMENT OF PATIENTS WITH CLL, FL, OR SLL

U-1616 NASAL ADMINISTRATION OF A TESTOSTERONE GEL TO A PATIENT TO TREAT THE PATIENT FOR A CONDITION ASSOCIATED WITH A DEFICIENCY OR ABSENCE OF ENDOGENOUS TESTOSTERONE

U-1617 METHOD OF TREATING MEDULLARY THYROID CANCER

U-1618 A METHOD OF TREATING A PATIENT SUFFERING FROM A PAIN ASSOCIATED SLEEP DISTURBANCE COMPRISING ADMINISTERING A LIQUID COMPOSITION FORMULATED INSIDE A SOFT GEL CAPSULE, AS CLAIMED, TO THE PATIENT

U-1619 TREATMENT OF IMMUNE (IDIOPATHIC) THROMBOCYTOPENIA (ITP)

U-1620 METHOD OF TREATMENT OF IRON-RELATED CONDITIONS WITH AT LEAST 0.6 GRAMS OF ELEMENTAL IRON VIA AN IRON CARBOHYDRATE COMPLEX, WITH A SUBSTANTIALLY NON-IMMUNOGENIC CARBOHYDRATE COMPONENT, IN ABOUT 15 MINUTES OR LESS

U-1621 PULMONARY ADMINISTRATION OF A COMPOSITION COMPRISING INSULIN BOUND TO A COMPLEXING AGENT.

U-1622 FOR THE TREATMENT OF POLYCYTHEMIA VERA

U-1623 USE OF EXENATIDE MAY RESULT IN REDUCTION IN APPETITE.

U-1624 TREATMENT OF UNRESECTABLE HEPATOCELLULAR CARCINOMA, ADVANCED RENAL CELL CARCINOMA, OR DIFFERENTIATED THYROID CARCINOMA.

U-1625 METHOD OF TREATING SCHIZOPHRENIA BY ADMINISTERING ILOPERIDONE TO A PATIENT BY REDUCING THE DOSE IN PATIENTS WHO ARE POOR METABOLIZERS OF CYP2D6

U-1626 A METHOD OF TREATING OR PREVENTING OCULAR PAIN AND BURNING

U-1627 TREATMENT OF ACUTE UNCOMPLICATED INFLUENZA IN ADULTS

U-1628 METHOD OF TREATING DISORDERS WITH AN ETIOLOGY COMPRISING OR ASSOCIATED WITH EXCESS GH-SECRETION

U-1629 METHOD OF TREATING ACROMEGALY

U-1630 TREATMENT IN COMBINATION WITH A CORTICOID SUCH AS PREDNISONE OF PROSTATE CANCER PREVIOUSLY TREATED WITH DOCETAXEL

U-1631 TREATMENT OF INFLAMMATORY LESIONS OF ROSACEA

U-1632 TREATMENT OF SCHIZOPHRENIA, WITH EFFICACY IN TREATING ACUTE EPISODES OF SCHIZOPHRENIA

U-1633 USE OF ARIPIPRAZOLE IN EXTENDED RELEASE INJECTABLE SUSPENSION IN TREATING ACUTE EPISODES OF SCHIZOPHRENIA

U-1634 TREATMENT OF BRCA MUTATED OVARIAN CANCER USING PARP INHIBITOR

U-1635 USE OF RITONAVIR AS A POTENT CYP3A INHIBITOR TO INCREASE PLASMA DRUG CONCENTRATION OF PARITAPREXIR AND OVERALL DRUG EXPOSURE FOR TREATMENT OF HCV INFECTION

U-1636 USE OF DASABUVIR TO INHIBIT VIRAL REPLICATION FOR THE TREATMENT OF HCV INFECTION.

U-1637 TREATMENT OF HCV INFECTION USING PARITAPREXIR, OMBITASVIR, RITONAVIR, AND DASABUVIR WITH RIBAVIRIN.

U-1638 TREATMENT OF HCV INFECTION USING PARITAPREXIR

U-1639 USE OF NALTREXONE AND BUPROPION IN EXTENDED-RELEASE FORM FOR CHRONIC WEIGHT MANAGEMENT FOR TREATING OVERWEIGHT OR OBESITY

U-1640 TREATMENT OF MODERATE TO SEVERE CHRONIC PAIN BY ADMINISTERING AN INTACT COMPOSITION AS CLAIMED

U-1641 MEMANTINE HCL/DONEPEZIL HCL COMBINATION FOR THE TREATMENT OF MODERATE TO SEVERE DEMENTIA OF THE ALZHEIMER'S TYPE

U-1642 METHOD OF TREATING TYPE 2 DIABETES MELLITUS IN PATIENTS WITH SEVERE CHRONIC RENAL IMPAIRMENT AND FOR WHOM METFORMIN THERAPY IS INAPPROPRIATE BY ADMINISTERING LINAGLIPTIN

U-1643 TREATING CUSHING'S SYNDROME

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-1644 TREATMENT OF OVERACTIVE BLADDER BY APPLICATION OF OXYBUTYNIN CHLORIDE GEL TO SKIN

U-1645 TREATMENT OF PARKINSON'S DISEASE, POST-ENCEPHALITIC PARKINSONISM, AND PARKINSONISM THAT MAY FOLLOW CARBON MONOXIDE INTOXICATION OR MANGANESE INTOXICATION

U-1646 TREATMENT OF POST-ENCEPHALITIC PARKINSONISM, AND PARKINSONISM THAT MAY FOLLOW CARBON MONOXIDE INTOXICATION OR MANGANESE INTOXICATION

U-1647 TREATMENT OF PARKINSONISM THAT MAY FOLLOW CARBON MONOXIDE INTOXICATION OR MANGANESE INTOXICATION

U-1648 TREATMENT OF PATIENTS WITH PARKINSON'S DISEASE, POST-ENCEPHALITIC PARKINSONISM, AND PARKINSONISM THAT MAY FOLLOW CARBON MONOXIDE INTOXICATION OR MANGANESE INTOXICATION

U-1649 TREATMENT OF POST-ENCEPHALITIC PARKINSONISM

U-1650 TREATMENT OF WALDENSTROM'S MACROGLOBULINEMIA

U-1651 METHOD OF TREATING TYPE 2 DIABETES MELLITUS BY ADMINISTERING LINAGLIPTIN IN COMBINATION WITH EMPAGLIFLOZIN

U-1652 METHOD OF TREATING TYPE 2 DIABETES MELLITUS BY ADMINISTERING LINAGLIPTIN IN COMBINATION WITH EMPAGLIFLOZIN AND METFORMIN

U-1653 METHOD OF TREATING TYPE 2 DIABETES MELLITUS BY ADMINISTERING LINAGLIPTIN IN COMBINATION WITH EMPAGLIFLOZIN (WITH OR WITHOUT METFORMIN)

U-1654 METHOD OF TREATING TYPE 2 DIABETES MELLITUS BY ADMINISTERING LINAGLIPTIN IN COMBINATION WITH EMPAGLIFLOZIN (WITH OR WITHOUT INSULIN OR A SULFONYLUREA)

U-1655 A METHOD TO ACCELERATE THE TIME TO GASTROINTESTINAL RECOVERY BY ADMINISTERING ABOUT 12 MG OF ALVIMOPAN TO THE PATIENT FROM ABOUT 30 TO 60 MINUTES PRIOR TO SURGERY

U-1656 METHOD OF IRON ADMINISTRATION TO TREAT PATIENTS IN NEED OF IRON REPLACEMENT

U-1657 METHOD FOR PROVIDING POST COITAL CONTRACEPTION TO A WOMAN BY ADMINISTERING ABOUT 30 MG OF ULIPRISTAL ACETATE WITHIN ABOUT 120 HOURS AFTER INTERCOURSE, WHEREIN THE WOMAN IS OVERWEIGHT HAVING A BMI OF 25 TO 29.99

U-1658 TREATMENT OF ER-POSITIVE, HER2-NEGATIVE ADVANCED BREAST CANCER IN COMBINATION WITH LETROZOLE AS INITIAL ENDOCRINE-BASED THERAPY FOR METASTATIC DISEASE IN POSTMENOPAUSAL WOMEN

U-1659 MANAGEMENT OF PAIN

U-1660 TREATMENT OF HIV-1 INFECTION IN ADULTS WITH NO DARUNAVIR RESISTANCE-ASSOCIATED SUBSTITUTIONS

U-1661 RISK-REDUCTION OF NSAID-ASSOCIATED GASTRIC ULCERS IN PATIENTS ALSO TAKING LOW DOSE ASPIRIN

U-1662 A METHOD OF TREATING OCULAR PAIN

U-1663 TREATMENT OF HIV-1 INFECTION

U-1664 TREATMENT OF BACTERIAL VAGINOSIS WITH METRONIDAZOLE GEL

U-1665 METHOD OF TREATING ATTENTION DEFICIT HYPERACTIVITY DISORDER BY ADMINISTERING THE COMPOSITION OF CLAIM 1

U-1666 PALLIATIVE TREATMENT OF PROSTATE CANCER

U-1667 TREATMENT OF ALLERGIC RHINITIS, INCLUDING SEASONAL ALLERGIC RHINITIS

U-1668 METHOD OF TREATING DEPRESSION OR MAJOR DEPRESSIVE DISORDER

U-1669 TREATMENT OF MULTIPLE MYELOMA, IN COMBINATION WITH BORTEZOMIB AND DEXAMETHASONE

U-1670 NATROBA TOPICAL SUSPENSION IS A PEDICULICIDE INDICATED FOR THE TOPICAL TREATMENT OF HEAD LICE INFESTATION IN PATIENTS SIX (6) MONTHS OF AGE AND OLDER.

U-1671 TREATMENT OF OCULAR ITCHING ASSOCIATED WITH CONJUNCTIVITIS

U-1672 TREATMENT OF COMPLICATED INTRA-ABDOMINAL INFECTION

U-1673 TREATMENT OF COMPLICATED URINARY TRACT INFECTION, INCLUDING PYELONEPHRITIS

U-1674 DOSAGE MODIFICATION TO REDUCE RISKS ASSOCIATED WITH QT PROLONGATION NOT INDUCED BY OTHER DRUGS DURING TREATMENT WITH ILOPERIDONE

U-1675 USE OF TROKENDI XR FOR THE TREATMENT OF EPILEPSY

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-1676 METHODS FOR TREATING BACTERIAL INFECTIONS

U-1677 TREATMENT OF IDIOPATHIC PULMONARY FIBROSIS (IPF)

U-1678 FOR THE TREATMENT OF PATIENTS WITH CLL, FL, OR SLL

U-1679 TREATMENT OF ACUTE OTITIS EXTERNA

U-1680 TREATMENT OF OCULAR ITCHING ASSOCIATED WITH ALLERGIC CONJUNCTIVITIS

U-1681 TREATMENT OF PATIENTS WITH PROGRESSIVE NEUROENDOCRINE TUMORS OF PANCREATIC ORIGIN (PNET) THAT ARE UNRESECTABLE, LOCALLY ADVANCED OR METASTATIC

U-1682 TREATMENT OF BACTERIAL VAGINOSIS

U-1683 TREATMENT FOR CHRONIC LYMPHOCYTIC LEUKEMIA WITH 17P DELETION

U-1684 TREATMENT OF CHRONIC LYMPHOCYTIC LEUKEMIA

U-1685 DOSAGE MODIFICATION TO REDUCE THE RISK ASSOCIATED WITH QT PROLONGATION NOT INDUCED BY OTHER DRUGS DURING TREATMENT WITH ILOPERIDONE

U-1686 A METHOD TO REDUCE WITHDRAWAL SYMPTOMS, INCLUDING NICOTINE CRAVING, ASSOCIATED WITH SMOKING CESSATION

U-1687 TREATMENT OF HCV INFECTION USING OMBITASVIR

U-1688 METHOD FOR CHRONIC WEIGHT MANAGEMENT BY TREATING OBESITY IN AN INDIVIDUAL WHO DOES NOT HAVE SEVERE RENAL IMPAIRMENT OR ESRD

U-1689 METHOD FOR CHRONIC WEIGHT MANAGEMENT BY INDUCING SATIETY IN AN INDIVIDUAL WHO DOES NOT HAVE SEVERE RENAL IMPAIRMENT OR ESRD

U-1690 METHOD FOR REDUCTION OF SUBMENTAL FAT

U-1691 INDICATED FOR THE ONCE-DAILY INHALED TREATMENT FOR ASTHMA IN ADULTS AGED 18 YEARS AND OLDER

U-1692 METHOD FOR CHRONIC WEIGHT MANAGEMENT BY DECREASING FOOD INTAKE IN AN INDIVIDUAL WHO DOES NOT HAVE SEVERE RENAL IMPAIRMENT OR ESRD

U-1693 METHOD OF TREATING ADHD IN CHILDREN 6 YEARS OF AGE AND OLDER AND ADOLESCENTS

U-1694 A METHOD FOR TREATING HEART FAILURE IN A HUMAN USING A CRYSTALLINE FORM OF IVABRADINE HYDROCHLORIDE

U-1695 METHOD FOR TREATING THYROID CARCINOMA INCLUDING DIFFERENTIATED THYROID CANCER

U-1696 TREATMENT OF UNRESECTABLE HEPATOCELLULAR CARCINOMA

U-1697 PULMONARY ADMINISTRATION OF A COMPOSITION COMPRISING INSULIN BOUND TO A DIKETOPIPERAZINE.

U-1698 PROPHYLAXIS OF INVASIVE ASPERGILLUS AND CANDIDA INFECTIONS

U-1699 A METHOD FOR TREATING ACUTE LYMPHOBLASTIC LEUKEMIA

U-1700 A METHOD FOR TREATING PHILADELPHIA CHROMOSOME POSITIVE ACUTE LYMPHOBLASTIC LEUKEMIA

U-1701 A METHOD FOR TREATING LEUKEMIA RESULTING FROM A MUTATION IN THE BCR-ABL KINASE DOMAIN

U-1702 TREATMENT OF COPD

U-1703 TREATMENT OF RESPIRATORY COMPLAINTS

U-1704 USE FOR TREATMENT IN PATIENTS WITH DIABETES

U-1705 USE FOR TREATMENT IN PATIENTS WITH HYPERGLYCEMIA

U-1706 TREATMENT OF TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE WHEREIN THE COMBINED THERAPEUTIC EFFECT IS GREATER THAN THE ADDITIVE EFFECT OF ADMINISTERING EACH AGENT ALONE

U-1707 TREATMENT OF IRRITABLE BOWEL SYNDROME WITH DIARRHEA (IBS-D) IN ADULTS AND SYMPTOMS THEREOF.

U-1708 TREATMENT OF IRRITABLE BOWEL SYNDROME WITH DIARRHEA (IBS-D) IN ADULTS.

U-1709 TREATMENT OF IRRITABLE BOWEL SYNDROME WITH DIARRHEA (IBS-D) WITH VIBERZI (ELUXADOLINE).

U-1710 TREATMENT OF NON-24-HOUR SLEEP-WAKE DISORDER BY AVOIDING THE USE OF TASIMELTEON IN COMBINATION WITH FLUVOXAMINE

U-1711 FOR THE TREATMENT OF PATIENTS WITH CLL, FL OR SLL

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

- U-1712 MEKINIST IN COMBINATION WITH DABRAFENIB FOR THE TREATMENT OF PATIENTS WITH UNRESECTABLE OR METASTATIC MELANOMA
- U-1713 TAFINLAR IN COMBINATION WITH TRAMETINIB FOR THE TREATMENT OF PATIENTS WITH UNRESECTABLE OR METASTATIC MELANOMA
- U-1714 TREATMENT OF THROMBOCYTOPENIA IN ADULT AND PEDIATRIC PATIENTS 6 YEARS AND OLDER WITH CHRONIC IMMUNE (IDIOPATHIC) THROMBOCYTOPENIA (ITP)
- U-1715 P2Y12 PLATELET INHIBITOR FOR USE AS ADJUNCT TO PERCUTANEOUS CORONARY INTERVENTION TO REDUCE RISK OF VARIOUS DISEASES/CONDITIONS IN PATIENTS NOT TREATED WITH A P2Y12 PLATELET INHIBITOR AND NOT GIVEN A GLYCOPROTEIN IIB/IIIA INHIBITOR
- U-1716 TREATMENT OF COUGH AND SYMPTOMS ASSOCIATED WITH UPPER RESPIRATORY ALLERGIES OR A COMMON COLD WITH CODEINE PHOSPHATE AND CHLORPHENIRAMINE MALEATE ORALLY ADMINISTERED EXTENDED RELEASE TABLETS
- U-1717 METHOD OF TREATING CYSTIC FIBROSIS IN PATIENTS WHO ARE HOMOZYGOUS FOR THE F508DEL MUTATION IN THE CYSTIC FIBROSIS TRANSMEMBRANE CONDUCTANCE REGULATOR (CFTR) GENE
- U-1718 METHOD OF TREATING CYSTIC FIBROSIS IN PATIENTS WHO HAVE THE F508DEL MUTATION IN THE CYSTIC FIBROSIS TRANSMEMBRANE CONDUCTANCE REGULATOR (CFTR) GENE.
- U-1719 ACUTE TREATMENT OF MIGRAINE
- U-1720 METHOD OF PROVIDING A THERAPEUTICALLY EFFECTIVE AND STABLE MEDIAN BLOOD PLASMA LEVEL OF LEVODOPA
- U-1721 USE OF RUXOLITINIB (JAKAFI) FOR BLOCKING SIGNAL TRANSDUCTION OF JANUS ASSOCIATED KINASES (JAKS) JAK1 AND/OR JAK2
- U-1722 TREATMENT OF BASAL CELL CARCINOMA
- U-1723 TREATMENT OF HEART FAILURE
- U-1724 METHOD OF INHIBITING HEPATITIS C VIRUS
- U-1725 METHOD OF INHIBITING HEPATITIS C VIRUS WITH DAKLINZA AND AT LEAST ONE ADDITIONAL COMPOUND HAVING ANTI-HCV ACTIVITY
- U-1726 REDUCTION IN RISK OF HOSPITALIZATION IN PATIENTS WITH CORONARY HEART DISEASE AND A HISTORY OF PAROXYSMAL OR PERSISTENT AF AND WITH ONE OR MORE RISK FACTORS BY ADMINISTRATION TWICE A DAY WITH MORNING AND EVENING MEALS
- U-1727 TOPICAL TREATMENT OF INFLAMMATORY PAPULES AND PUSTULES OF MILD TO MODERATE ROSACEA
- U-1728 REDUCTION IN RISK OF HOSPITALIZATION IN PATIENTS WITH STABLE NYHA CLASS III HEART FAILURE AND A HISTORY OF PAROXYSMAL OR PERSISTENT AF AND WITH ONE OR MORE RISK FACTORS BY ADMINISTRATION TWICE A DAY WITH MORNING AND EVENING MEALS
- U-1729 REDUCE THE RISK OF RECURRENT DEEP VEIN THROMBOSIS (DVT)
- U-1730 REDUCE THE RISK OF RECURRENT PULMONARY EMBOLISM
- U-1731 TEMPORARY RELIEF OF MINOR ACHES AND PAINS
- U-1732 TEMPORARY REDUCTION OF FEVER
- U-1733 TREATMENT/PREVENTION OF CARDIOVASCULAR DISEASE
- U-1734 USE OF FLIBANSERIN OR A PHARMACEUTICALLY ACCEPTABLE ACID ADDITION SALT THEREOF TO TREAT HYPOACTIVE SEXUAL DESIRE DISORDER (HSDD)
- U-1735 METHODS OF TREATING PAIN, INFLAMMATION AND/OR FEVER WITH INTRAVENOUS IBUPROFEN SUCH THAT MEAN ARTERIAL BLOOD PRESSURE DOES NOT INCREASE THE DOSAGE INTERVAL
- U-1736 TREATMENT OF THROMBOCYTOPENIA IN ADULT AND PEDIATRIC PATIENTS 1 YEAR AND OLDER WITH CHRONIC IMMUNE (IDIOPATHIC) THROMBOCYTOPENIA (ITP)
- U-1737 METHOD OF TREATING SCHIZOPHRENIA BY ADMINISTERING ILOPERIDONE TO A PATIENT BY REDUCING THE DOSE IN PATIENTS WHO ARE BEING TREATED WITH FLUOXETINE
- U-1738 TREATMENT OF IRRITABLE BOWEL SYNDROME WITH DIARRHEA (IBS-D) WITH VIBERZI (ELUXADOLINE)
- U-1739 MANAGEMENT OF PAIN SEVERE ENOUGH TO REQUIRE DAILY, AROUND -THE-CLOCK, LONG-TERM OPIOID TREATMENT, INCLUDING NEUROPATHIC PAIN ASSOCIATED WITH DIABETIC PERIPHERAL NEUROPATHY
- U-1740 IN COMBINATION WITH OTHER ANTIRETROVIRAL AGENTS FOR THE TREATMENT OF HIV-1 INFECTION IN TREATMENT-NAIVE PATIENTS WITH HIV-1 RNA LESS THAN OR EQUAL TO 100,000 AT THE START OF THERAPY

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-1741 PREVENTION OF DELAYED NAUSEA AND VOMITING ASSOCIATED WITH EMETOGENIC CANCER CHEMOTHERAPY

U-1742 ROLAPITANT IS APPROVED FOR THE PREVENTION OF DELAYED NAUSEA AND VOMITING (I.E., EMESIS) ASSOCIATED WITH EMETOGENIC CANCER CHEMOTHERAPY

U-1743 FOR THE PREVENTION OF NAUSEA AND VOMITING ASSOCIATED WITH CHEMOTHERAPY

U-1744 PREVENTION OF POST-OPERATIVE NAUSEA AND VOMITING

U-1745 FOR THE TREATMENT OF PATIENTS WITH WALDENSTROM'S MACROGLOBULINEMIA

U-1746 MONOTHERAPY OR ADJUNCTIVE THERAPY FOR TREATMENT OF PARTIAL-ONSET SEIZURES AND APPROVED IN PATIENTS WITH EPILEPSY

U-1747 FOR CLAIMS 1-3,6-13,16-24 AND 26-32: METHOD OF TREATING ADHD

U-1748 FOR CLAIMS 1-4,6-14,16-24 AND 26-32: METHOD OF TREATING ADHD IN CHILDREN 6 YEARS OF AGE AND OLDER AND ADOLESCENTS

U-1749 ACUTE TREATMENT OF MANIC AND MIXED EPISODES ASSOCIATED WITH BIPOLAR I DISORDER

U-1750 TREATMENT OF SCHIZOPHRENIA AND/OR ACUTE MANIC OR MIXED EPISODES ASSOCIATED WITH BIPOLAR I DISORDER WITH CARIPRAZINE

U-1751 TREATMENT OF PATIENTS WITH METASTATIC COLORECTAL CANCER WHO HAVE BEEN PREVIOUSLY TREATED WITH FLUOROPYRIMIDINE-, OXALIPLATIN- AND IRINOTECAN-BASED CHEMOTHERAPY, AN ANTI-VEGF BIOLOGICAL THERAPY, AND IF RAS WILD-TYPE, AN ANTI-EGFR THERAPY

U-1752 PROPHYLAXIS OF ORGAN REJECTION

U-1753 TREATMENT OF HCV INFECTION USING DASABUVIR

U-1754 FOR THE TREATMENT OF PULMONARY HYPERTENSION (PAH) IN COMBINATION WITH TADALAFIL

U-1755 FIRST-LINE TREATMENT OF PATIENTS WITH METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) WHOSE TUMORS HAVE EPIDERMAL GROWTH FACTOR RECEPTOR (EGFR) EXON 19 DELETIONS OR EXON 21 (L858R) SUBSTITUTION MUTATIONS

U-1756 METHODS OF TREATING PAIN, INFLAMMATION AND/OR FEVER IN A CRITICALLY ILL PATIENT WITH INTRAVENOUS IBUPROFEN IN NEED THEREOF

U-1757 INHIBITION ON PI3K KINASE

U-1758 METHOD OF TREATING ALLERGIC REACTION VIA INJECTION

U-1759 METHOD OF REVERSING THE ANTICOAGULANT EFFECT OF DABIGATRAN USING IDARUCIZUMAB

U-1760 RISK-REDUCTION OF NSAID GASTRIC ULCER IN PATIENTS REQUIRING CHRONIC NSAID TREATMENT

U-1761 PLAQUE PSORIASIS

U-1762 USE OF BELVIQ (LORCASERIN HYDROCHLORIDE) FOR CHRONIC WEIGHT MANAGEMENT IN PATIENTS ON A REDUCED-CALORIE DIET AND WHO HAVE ACHIEVED A GREATER THAN OR EQUAL TO 5% WEIGHT LOSS BY WEEK 12 OF TREATMENT

U-1763 USE OF BELVIQ (LORCASERIN HYDROCHLORIDE) FOR CHRONIC WEIGHT MANAGEMENT BY DECREASING FOOD INTAKE IN PATIENTS ON A REDUCED-CALORIE DIET AND WHO HAVE ACHIEVED GREATER THAN OR EQUAL TO 5% WEIGHT LOSS BY WEEK 12 OF TREATMENT

U-1764 USE OF BELVIQ (LORCASERIN HYDROCHLORIDE) FOR CHRONIC WEIGHT MANAGEMENT BY INDUCING SATIETY IN PATIENTS ON A REDUCED-CALORIE DIET AND WHO HAVE ACHIEVED A GREATER THAN OR EQUAL TO 5% WEIGHT LOSS BY WEEK 12 OF TREATMENT

U-1765 USE OF BELVIQ (LORCASERIN HYDROCHLORIDE) FOR CHRONIC WEIGHT MANAGEMENT BY TREATING OBESITY IN PATIENTS ON A REDUCED-CALORIE DIET AND WHO HAVE ACHIEVED A GREATER THAN OR EQUAL TO 5% WEIGHT LOSS BY WEEK 12 OF TREATMENT

U-1766 TREATMENT OF HYPERKALEMIA

U-1767 USE OF CALCIPOTRIENE FOAM FOR THE TOPICAL TREATMENT OF PLAQUE PSORIASIS IN PATIENTS AGED 18 YEARS AND OLDER

U-1768 METHOD OF TREATING TYPE 2 DIABETES MELLITUS BY ADMINISTERING LINAGLIPTIN

U-1769 TREATMENT OF PAIN BY TRANSMUCOSAL DELIVERY OF BUPRENORPHINE

U-1770 TREATMENT OF SCHIZOPHRENIA WITH IMPROVEMENT IN NEGATIVE SYMPTOMS AND/OR COGNITIVE DYSFUNCTION OF SCHIZOPHRENIA

U-1771 ADMINISTRATION OF REMODULIN DILUTED FOR INTRAVENOUS INFUSION WITH STERILE WATER FOR INJECTION OR 0.9% SODIUM CHLORIDE INJECTION PRIOR TO ADMINISTRATION

U-1772 METHOD OF TREATING TYPE 2 DIABETES MELLITUS BY ADMINISTERING LINAGLIPTIN IN COMBINATION WITH EMPAGLIFLOZIN

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-1773 LONG-TERM MAINTENANCE TREATMENT OF AIRFLOW OBSTRUCTION IN PATIENTS WITH CHRONIC OBSTRUCTIVE PULMONARY DISEASE (COPD)

U-1774 USE OF A LOTION CONTAINING HALOBETASOL PROPIONATE FOR THE TREATMENT OF CORTICOSTEROID-RESPONSIVE

U-1775 USE OF A LOTION CONTAINING HALOBETASOL PROPIONATE FOR THE TREATMENT OF CORTICOSTEROID-RESPONSIVE DERMATOSES INCLUDING PSORIASIS

U-1776 METHOD OF USING COBIMETINIB FOR THE TREATMENT OF MELANOMA

U-1777 TREATMENT OF PATIENTS WITH METASTATIC EPIDERMAL GROWTH FACTOR RECEPTOR (EGFR) T790M MUTATION-POSITIVE NON-SMALL CELL LUNG CANCER (NSCLC), WHO HAVE PROGRESSED ON OR AFTER EGFR TKI THERAPY

U-1778 METHOD FOR TREATING MULTIPLE MYELOMA

U-1779 METHOD FOR TREATING MULTIPLE MYELOMA WITH ONE OR MORE OTHER THERAPEUTIC AGENTS

U-1780 METHOD FOR TREATING CANCER, INCLUDING MULTIPLE MYELOMA

U-1781 RISK-REDUCTION OF NSAID-ASSOCIATED GASTRIC ULCER IN PATIENTS REQUIRING NSAID TREATMENT

U-1782 FOR HEAD LICE INFESTATIONS

U-1783 METHOD OF TREATING FREQUENT HEARTBURN BY ADMINISTERING AN ESOMEPRAZOLE MAGNESIUM AS CLAIMED

U-1784 METHOD OF TREATING FREQUENT HEARTBURN BY ADMINISTERING AN ESOMEPRAZOLE MAGNESIUM TRIHYDRATE AS CLAIMED

U-1785 METHOD OF TREATING FREQUENT HEARTBURN BY ADMINISTERING AN ESOMEPRAZOLE MAGNESIUM FORMULATION AS CLAIMED

U-1786 TREATMENT OF PATIENTS WITH RELAPSING FORMS OF MULTIPLE SCLEROSIS WHILE MANAGING THE RISK OF TERIFLUNOMIDE AND ROSUVASTATIN INTERACTION BY LIMITING THE ROSUVASTATIN DOSE TO NO MORE THAN 10MG AND/OR ADMINISTERING ABOUT HALF THE NORMAL DOSE

U-1787 TREATMENT OF EXOCRINE PANCREATIC INSUFFICIENCY

U-1788 TREATMENT OF PATIENT HAVING DIABETES MELLITUS VIA ORAL INHALATION OF FDKP MICROPARTICLES COMPRISING INSULIN

U-1789 METHOD OF ADMINISTERING AN ETHANOL-FREE TAXANE LIQUID NANODISPERSION FORMULATION TO A SUBJECT COMBINING THE FORMULATION WITH AN AQUEOUS MEDIUM TO PROVIDE AN ETHANOL-FREE TAXANE DILUTED SOLUTION

U-1790 FOR USE IN TREATMENT OF PATIENTS WITH CHRONIC LYMPHOCYTIC LEUKEMIA (CLL) AND/OR NON-HODGKIN'S LYMPHOMA

U-1791 EMERGENCY TREATMENT OF ADULT & PEDIATRIC PATIENTS FOLLOWING FLUOROURACIL OR CAPECITABINE OVERDOSE, OR WHO EXHIBIT EARLY-ONSET, SEVERE OR LIFE-THREATENING CARDIAC OR CNS TOXICITY OR UNUSUALLY SEVERE ADVERSE REACTIONS WITHIN 96 HOURS

U-1792 TREATMENT OF OTIC INFECTION OR INFLAMMATION

U-1793 TREATMENT OF PEDIATRIC PATIENTS WITH OTITIS MEDIA WITH EFFUSION UNDERGOING TYMPANOSTOMY TUBE PLACEMENT

U-1794 REVERSAL OF DRUG-INDUCED NEUROMUSCULAR BLOCK

U-1795 REVERSAL OF NEUROMUSCULAR BLOCKAGE INDUCED BY ROCURONIUM BROMIDE OR VECURONIUM BROMIDE

U-1796 TOPICAL TREATMENT OF INFLAMMATORY PAPULES AND PUSTULES OF MILD TO MODERATE ROSACEA

U-1797 METHOD OF TREATING PULMONARY ARTERIAL HYPERTENSION COMPRISING ADMINISTERING A PHARMACEUTICAL COMPOSITION COMPRISING SELEXIPAG

U-1798 METHOD OF TREATING PULMONARY ARTERIAL HYPERTENSION COMPRISING ADMINISTERING SELEXIPAG IN COMBINATION WITH THE ENDOTHELIN RECEPTOR ANTAGONIST MACITENTAN

U-1799 METHOD OF INCREASING GROWTH OF HAIR INCLUDING EYELASHES

U-1800 A METHOD OF TREATING OCULAR PAIN AND/OR ENHANCING OCULAR COMFORT

U-1801 REDUCTION OF SERUM URIC ACID LEVELS

U-1802 TREATMENT OF GOUT

U-1803 TREATMENT OF HYPERURICEMIA

U-1804 ACHIEVING A THERAPEUTIC BENEFIT IN A SUBJECT WITH GOUT

PATENT AND EXCLUSIVITY TERMS

ADB 131 of 225

PATENT USE

- U-1805 USE OF DEXLANSOPRAZOLE IN PATIENTS TAKING CLOPIDOGREL WITHOUT MEANINGFUL CYP2C19 INTERACTIONS
- U-1806 COADMINISTERING WITH ALLOPURINOL TO REDUCE SERUM URIC ACID (SUA) BELOW 4 MG/DL; BELOW 6MG/DL IN PATIENTS HAVING URIC ACID DEPOSITS; AND/OR BELOW 6MG/DL WITH SUA INTRADAY CHANGE MORE THAN 50% AND/OR ADVERSE EVENT RATE LESS THAN 15%
- U-1807 TREATMENT OF PEDIATRIC PATIENTS 8 TO 17 YEARS OF AGE WITH HETEROZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA (HEFH)
- U-1808 USE OF NALTREXONE AND BUPROPION FOR CHRONIC WEIGHT MANAGEMENT FOR TREATING OVERWEIGHT OR OBESITY IN PATIENTS WITH MAJOR DEPRESSIVE DISORDER
- U-1809 METHOD OF DRUG DELIVERY VIA THE NASAL CAVITY
- U-1810 TREATMENT OF PAIN IN PATIENTS WITH HEPATIC IMPAIRMENT
- U-1811 TREATMENT OF PATIENTS WITH UNRESECTABLE OR METASTATIC MELANOMA WITH BRAF V600E MUTATIONS AFTER CONFIRMING THE PRESENCE OF BRAF V600E MUTATION
- U-1812 TREATMENT OF PATIENTS WITH UNRESECTABLE OR METASTATIC LIPOSARCOMA
- U-1813 TREATMENT OF PATIENTS INFECTED WITH HEPATITIS C VIRUS
- U-1814 METHOD OF TREATING GLAUCOMA OR ELEVATED INTRAOCULAR PRESSURE
- U-1815 TREATMENT OF PARTIAL-ONSET SEIZURES AS ADJUNCTIVE THERAPY IN PATIENTS WITH EPILEPSY AGED 16 YEARS AND OLDER WITH EPILEPSY
- U-1816 TREATMENT OF A UREA CYCLE DISORDER
- U-1817 PEDIATRIC USE AGES 1 MONTH TO 2 YEARS, GERD AND EROSIIVE ESOPHAGITIS
- U-1818 TREATING HR-POS., HER2-NEG. ADVANCED OR METASTATIC BREAST CANCER WITH PALBOCICLIB IN COMBO WITH LETROZOLE AS INITIAL ENDOCRINE BASED THERAPY IN POSTMENOPAUSAL WOMEN, OR FULVESTRANT IN WOMEN WITH DISEASE PROGRESSION AFTER ENDOCRINE THERAPY
- U-1819 MANAGEMENT OF PAIN SEVERE ENOUGH TO REQUIRE DAILY, AROUND-THE-CLOCK, LONG TERM OPIOID TREATMENT AND FOR WHICH ALTERNATIVE TREATMENT OPTIONS ARE INADEQUATE
- U-1820 METHOD OF TREATING PULMONARY ARTERIAL HYPERTENSION BY ADMINISTERING A PHARMACEUTICAL COMPOSITION COMPRISING MACITENTAN AND A POLYSORBATE, WHEREIN THE POLYSORBATE REPRESENTS 0.1 TO 3% OF THE WEIGHT OF SAID PHARMACEUTICAL COMPOSITION
- U-1821 METHOD FOR CONTRACEPTION TO A WOMAN COMPRISING ADMINISTERING TO THE WOMAN 30MG OF ULIPRISTAL ACETATE MORE THAN 72 HOURS AND UP TO 120 HOURS AFTER AN UNPROTECTED INTERCOURSE
- U-1822 TREATMENT OF SCHIZOPHRENIA OR BIPOLAR DEPRESSION WITH IMPROVEMENT IN ATTENTION FUNCTION IN SCHIZOPHRENIA AND/OR BIPOLAR DISORDER
- U-1823 A METHOD OF PROVIDING NITRIC OXIDE THERAPY TO A PATIENT BY COMPENSATING LONG-TERM SENSITIVITY DRIFT OF ELECTROCHEMICAL GAS SENSORS USED IN SYSTEMS FOR DELIVERING THERAPEUTIC NITRIC OXIDE TO A PATIENT
- U-1824 A METHOD OF PROVIDING NITRIC OXIDE THERAPY TO A PATIENT BY VERIFYING GAS INFORMATION OF NITRIC OXIDE PRIOR TO DELIVERY TO PATIENT
- U-1825 METHOD OF USING VISMODEGIB TO TREAT CANCER IN A MAMMAL
- U-1826 TREATMENT OF HR-POSITIVE, HUMAN EPIDERMAL GROWTH FACTOR RECEPTOR 2 (HER2)-NEGATIVE ADVANCED OR METASTATIC BREAST CANCER IN COMBINATION WITH PALBOCICLIB IN WOMEN WITH DISEASE PROGRESSION AFTER ENDOCRINE THERAPY
- U-1827 A METHOD OF PROVIDING A SUBJECT WITH THERAPEUTICALLY EFFECTIVE AMOUNT OF RACEMIC METHYLPHENIDATE BY ORALLY ADMINISTERING TO SAID SUBJECT A SINGLE METHYLPHENIDATE EXTENDED RELEASE CHEWABLE TABLET ACCORDING TO CLAIM 1
- U-1828 INCREASING MEAN ARTERIAL BLOOD PRESSURE IN ADULT PATIENTS WITH HYPOTENSION ASSOCIATED WITH SEPTIC SHOCK
- U-1829 EMERGENCY TREATMENT OF ALLERGIC REACTIONS (TYPE I), INCLUDING ANAPHYLAXIS
- U-1830 INDUCTION AND MAINTENANCE OF MYDRIASIS DURING INTRAOCULAR SURGERY
- U-1831 METHOD OF TREATING PULMONARY ARTERIAL HYPERTENSION COMPRISING ADMINISTERING A CRYSTALLINE FORM OF SELEXIPAG
- U-1832 IMPROVEMENT IN GLYCEMIC CONTROL IN DIABETES MELLITUS PATIENTS BY USE OF A PEN INJECTOR WITH A THREADED DRIVE SLEEVE
- U-1833 REDUCTION OF ELEVATED INTRAOCULAR PRESSURE IN PATIENTS WITH OPEN ANGLE GLAUCOMA

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

OR OCULAR HYPERTENSION

- U-1834 TREATMENT OF POSTOPERATIVE INFLAMMATION AND PREVENTION OF OCULAR PAIN IN PATIENTS UNDERGOING CATARACT SURGERY
- U-1835 TREATMENT OF PATIENTS WITH CHRONIC LYMPHOCYTIC LEUKEMIA (CLL) WITH 17P DELETION, AS DETECTED BY AN FDA APPROVED TEST, WHO HAVE RECEIVED AT LEAST ONE PRIOR THERAPY
- U-1836 TREATMENT OF HEREDITARY TYROSINEMIA TYPE 1 (HT-1) IN COMBINATION WITH DIETARY RESTRICTION OF TYROSINE AND PHENYLALANINE
- U-1837 METHOD FOR TREATING TYPE II DIABETES MELLITUS BY ADMINISTERING SAXAGLIPTIN ALONE OR IN COMBINATION WITH INSULIN, METFORMIN, A THIAZOLIDINEDIONE, GLYBURIDE OR METFORMIN PLUS A SULFONYLUREA
- U-1838 METHOD FOR TREATING TYPE II DIABETES MELLITUS BY ADMINISTERING SAXAGLIPTIN IN COMBINATION WITH METFORMIN
- U-1839 COMPOSITION AND METHOD FOR PROVIDING A REDUCTION IN SIDE EFFECTS FOR HUMAN PATIENTS IN NEED OF ACETYLCYSTEINE THERAPY
- U-1840 TREATMENT OF HCV INFECTION USING PARITAPREXVIR, OMBITASVIR, RITONAVIR, AND DASABUVIR, WITHOUT RIBAVIRIN
- U-1841 USE IN THE LONG-TERM, MAINTENANCE TREATMENT OF AIRFLOW OBSTRUCTION IN PATIENTS WITH CHRONIC OBSTRUCTIVE PULMONARY DISEASE (COPD)
- U-1842 METHOD OF TREATING EPILEPSY
- U-1843 TREATMENT OF PSYCHOSIS
- U-1844 TREATMENT OF PARKINSON'S DISEASE PSYCHOSIS
- U-1845 TREATMENT OF PSYCHOSIS OR A SYMPTOM THEREOF
- U-1846 TREATMENT OF A NEURODEGENERATIVE DISEASE OR A SYMPTOM THEREOF
- U-1847 METHOD OF TREATING A BACTERIAL INFECTION
- U-1848 TREATMENT OF METASTATIC ADENOCARCINOMA OF THE PANCREAS THAT HAS PROGRESSED ON GEMCITABINE-BASED THERAPY, IN COMBINATION WITH 5-FLUOROURACIL AND LEUCOVORIN
- U-1849 METHOD OF TREATING PULMONARY HYPERTENSION BY ADMINISTERING TREPROSTINIL OR A SALT THEREOF BY INHALATION USING A DEVICE
- U-1850 METHOD OF ADMINISTERING LEVETIRACETAM
- U-1851 A DOSING REGIMEN FOR THE TREATMENT OF HYPERCHOLESTEROLEMIA AND HYPERLIPIDEMIA IN PATIENTS WITH HOMOZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA USING AT LEAST THREE STEP-WISE INCREASING DOSES
- U-1852 METHOD OF TREATING TYPE 2 DIABETES
- U-1853 METHOD OF TREATING TYPE 2 DIABETES MELLITUS BY ADMINISTERING A DIPEPTIDYL PEPTIDASE-IV INHIBITOR IN COMBINATION WITH METFORMIN AND, OPTIONALLY, A SULFONYLUREA
- U-1854 TREATMENT OF PRIMARY BILIARY CHOLANGITIS (PBC)
- U-1855 IMPROVEMENT IN GLYCEMIC CONTROL IN DIABETES MELLITUS PATIENTS
- U-1856 TREATMENT OF METASTATIC ADENOCARCINOMA OF THE PANCREAS THAT HAS PROGRESSED ON GEMCITABINE-BASED THERAPY, IN COMBINATION WITH 5-FLUOROURACIL AND LEUCOVORIN, IN A PATIENT HOMOZYGOUS FOR THE UGT1A1*28 ALLELE
- U-1857 TO INCREASE BLOOD PRESSURE IN ADULTS WITH VASODILATORY SHOCK (E.G., POST-CARDIOTOMY OR SEPSIS) WHO REMAIN HYPOTENSIVE DESPITE FLUIDS AND CATECHOLAMINES
- U-1858 TREATMENT OF PLAQUE PSORIASIS
- U-1859 TREATMENT OF SCHIZOPHRENIA, ACUTE TREATMENT OF MANIC AND MIXED EPISODES ASSOCIATED WITH BIPOLAR I DISORDER, ADJUNCTIVE TREATMENT OF MAJOR DEPRESSIVE DISORDER, AND TREATMENT OF IRRITABILITY ASSOCIATED WITH AUTISTIC DISORDER
- U-1860 REDUCTION OF THE RATE OF CARDIOVASCULAR DEATH, MYOCARDIAL INFARCTION, AND STROKE IN PATIENTS WITH ACUTE CORONARY SYNDROME OR A HISTORY OF MYOCARDIAL INFARCTION
- U-1861 USE OF AN INHALER TO ADMINISTER DRY POWDER MEDICAMENT
- U-1862 TREATMENT OF POST-MYOCARDIAL INFARCTION
- U-1863 TREATMENT OF STROKE
- U-1864 TREATMENT OF MYOCARDIAL INFARCTION

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-1865 TREATMENT OF THROMBOTIC STROKE

U-1866 TREATMENT OF STABLE AND UNSTABLE ANGINA

U-1867 METHOD OF INHIBITING PLATELET AGGREGATION

U-1868 TREATMENT OF ARTERIAL THROMBOTIC COMPLICATIONS SELECTED FROM THE GROUP CONSISTING OF UNSTABLE ANGINA, THROMBOTIC OR EMBOLIC STROKE, TRANSIENT ISCHAEMIC ATTACKS, PERIPHERAL VASCULAR DISEASE AND MYOCARDIAL INFARCTION

U-1869 TREATMENT OF AN ARTERIAL THROMBOTIC COMPLICATION IN A PATIENT WITH CORONARY ARTERY, CEREBROVASCULAR OR PERIPHERAL VASCULAR DISEASE

U-1870 ZINGO IS A POWDER INTRADERMAL SYSTEM THAT IS CAPABLE OF DELIVERING FINE DRY POWDERED LIDOCAINE HYDROCHLORIDE MONOHYDRATE FOR LOCAL ANESTHETIC ACTION

U-1871 TREATMENT OF SECONDARY HYPERPARATHYROIDISM IN PATIENTS WITH STAGE 3 OR 4 CHRONIC KIDNEY DISEASE USING CONTROLLED RELEASE, ORAL 25-HYDROXYVITAMIN D

U-1872 USE OF SUSTAINED RELEASE 25-HYDROXYVITAMIN D IN TREATING PATIENTS HAVING 25-HYDROXYVITAMIN D INSUFFICIENCY OR DEFICIENCY

U-1873 ADMINISTRATION OF 25-HYDROXYVITAMIN D3 BY CONTROLLED RELEASE

U-1874 TREATMENT OF FREQUENT HEARTBURN BY ADMINISTERING OMEPRAZOLE ACCORDING TO CLAIMS 1-8

U-1875 TREATMENT OF FREQUENT HEARTBURN BY ADMINISTERING S-OMEPRAZOLE TRIHYDRATE ACCORDING TO CLAIMS 1-3

U-1876 METHOD OF ANESTHETIZING AT LEAST A PORTION OF THE MAXILLARY DENTAL ARCH

U-1877 METHOD OF TREATING PULMONARY HYPERTENSION BY ORALLY ADMINISTERING A FORMULATION OF A PHARMACEUTICALLY ACCEPTABLE SALT OF TREPROSTINIL

U-1878 FOR OPIOID DEPENDENCE

U-1879 METHOD OF DIAGNOSING TUMORS USING POSITRON EMISSION TOMOGRAPHY

U-1880 TREATMENT OF SIGNS AND SYMPTOMS OF DRY EYE DISEASE (DED)

U-1881 IMPROVEMENT IN GLYCEMIC CONTROL IN TYPE 2 DIABETES MELLITUS PATIENTS BY USE OF A PEN INJECTOR

U-1882 MANAGEMENT OF MILD TO MODERATE PAIN, MANAGEMENT OF MODERATE TO SEVERE PAIN AS AN ADJUNCT TO OPIOID ANALGESICS, REDUCTION IN FEVER THROUGH ANALGESIC AND ANTIPYRETIC ACTIVITY

U-1883 TREATMENT OF GASTROINTESTINAL STROMAL TUMORS (GIST)

U-1884 USE OF LORCASERIN HYDROCHLORIDE FOR CHRONIC WEIGHT MANAGEMENT IN PATIENTS ON A REDUCED-CALORIE DIET AND WHO HAVE ACHIEVED A GREATER THAN OR EQUAL TO 5% WEIGHT LOSS BY WEEK 12 OF TREATMENT

U-1885 USE OF LORCASERIN HYDROCHLORIDE FOR CHRONIC WEIGHT MANAGEMENT BY DECREASING FOOD INTAKE IN PATIENTS ON A REDUCED-CALORIE DIET AND WHO HAVE ACHIEVED A GREATER THAN OR EQUAL TO 5% WEIGHT LOSS BY WEEK 12 OF TREATMENT

U-1886 USE OF LORCASERIN HYDROCHLORIDE FOR CHRONIC WEIGHT MANAGEMENT BY INDUCING SATIETY IN PATIENTS ON A REDUCED-CALORIE DIET AND WHO HAVE ACHIEVED A GREATER THAN OR EQUAL TO 5% WEIGHT LOSS BY WEEK 12 OF TREATMENT

U-1887 USE OF LORCASERIN HYDROCHLORIDE FOR CHRONIC WEIGHT MANAGEMENT BY TREATING OBESITY IN PATIENTS ON A REDUCED-CALORIE DIET AND WHO HAVE ACHIEVED A GREATER THAN OR EQUAL TO 5% WEIGHT LOSS BY WEEK 12 OF TREATMENT

U-1888 USE OF CONTROLLED RELEASE 25-HYDROXYVITAMIN D IN TREATING SECONDARY HYPERPARATHYROIDISM IN PATIENTS HAVING CHRONIC KIDNEY DISEASE

U-1889 TREATMENT OF HCV INFECTION USING DASABUVIR/OMBITASVIR/PARITAPREVIR/RITONAVIR FIXED DOSE COMBINATION

U-1890 OTC USE: ALLERGY SYMPTOM RELIEVER; TEMPORARILY RELIEVES THESE SYMPTOMS DUE TO HAY FEVER OR OTHER UPPER RESPIRATORY ALLERGIES; NASAL CONGESTION, RUNNY NOSE, SNEEZING, ITCHY NOSE, AND (ITCHY WATER EYES (AGES 12 AND UP))

U-1891 TREATMENT OR PREVENTION OF NAUSEA AND VOMITING

U-1892 METHOD OF TREATING LEFT VENTRICULAR DYSFUNCTION

U-1893 METHOD OF TREATING MANIC OR MIXED EPISODES ASSOCIATED WITH BIPOLAR DISORDER IN PEDIATRIC PATIENTS

U-1894 COMBINATION TREATMENT WITH A GLITAZONE FOR IMPROVEMENT OF GLYCEMIC CONTROL IN TYPE 2 DIABETES MELLITUS PATIENTS

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-1895 METHOD OF TREATING PROSTATE CANCER

U-1896 SUPPLEMENT FOR VITAMIN B12 DEFICIENCIES

U-1897 METHOD OF TREATING ACS USING ANGIOPLASTY WITH AGGRASTAT (TIROFIBAN HYDROCHLORIDE)

U-1898 METHOD OF INHIBITING PLATELET AGGREGATION WITH AGGRASTAT (TIROFIBAN HYDROCHLORIDE)

U-1899 TREATMENT OF PANCREATIC CANCER THAT HAS PROGRESSED ON GEMCITABINE-BASED THERAPY, IN COMBINATION WITH 5-FLUOROURACIL AND LEUCOVORIN

U-1900 TREATMENT OF THE SIGNS AND SYMPTOMS OF DRY EYE DISEASE (DED)

U-1901 TREATMENT OF SCHIZOAFFECTIVE DISORDER AS A MONOTHERAPY AND AS AN ADJUNCT TO MOOD STABILIZERS OR ANTIDEPRESSANTS

U-1902 TREATMENT OR SECONDARY PREVENTION OF CARDIOVASCULAR DISEASE, CARDIOVASCULAR EVENTS, OR CEREBROVASCULAR EVENTS AND RISK-REDUCTION OF ASPIRIN-ASSOCIATED GASTRIC ULCERS

U-1903 USE OF NALOXONE HYDROCHLORIDE FOR EMERGENCY TREATMENT OF KNOWN OR SUSPECTED OPIOID OVERDOSE, AS MANIFESTED BY RESPIRATORY AND/OR CENTRAL NERVOUS SYSTEM DEPRESSION.

U-1904 (I)TREATMENT OF DUCHENNE MUSCULAR DYSTROPHY; (II)RESTORING/INCREASING FUNCTIONAL DYSTROPHIN PROTEIN; OR (III) INDUCING SKIPPING; EACH OF (I)-(III) IN PATIENTS HAVING A CONFIRMED MUTATION OF THE DMD GENE THAT IS AMENABLE TO EXON 51 SKIPPING

U-1905 METHOD OF TREATING A PATIENT HAVING CYSTIC FIBROSIS, THE PATIENT HAVING A R117H MUTATION IN CFTR, USING N-(5-HYDROXY-2,4-DI-TERT-BUTYL-PHENYL)-4-OXO-1H-QUINOLINE-3-CARBOXAMIDE

U-1906 METHOD OF TREATING A PATIENT HAVING CYSTIC FIBROSIS, SUCH AS A PATIENT HAVING A G551D MUTATION IN CFTR, USING N-(5-HYDROXY-2,4-DI-TERT-BUTYL-PHENYL)-4-OXO-1H-QUINOLINE-3-CARBOXAMIDE

U-1907 USE OF A DELIVERY DEVICE TO ADMINISTER A DOSE OF NALOXONE

U-1908 METHOD OF TREATING CYSTIC FIBROSIS IN A PATIENT, THE PATIENT HAVING THE F508DEL MUTATION IN CFTR, USING IVACAFTOR AND FORM I LUMACAFTOR

U-1909 METHOD OF TREATING CYSTIC FIBROSIS IN A PATIENT, THE PATIENT HAVING THE F508DEL MUTATION IN CFTR, USING IVACAFTOR AND LUMACAFTOR

U-1910 METHOD OF TREATING CYSTIC FIBROSIS IN A PATIENT, THE PATIENT HAVING THE F508DEL MUTATION IN CFTR, USING THE DOSAGE UNIT OF CLAIM 1 OF U.S. PATENT NO. 8,716,338

U-1911 METHOD OF TREATING A PATIENT HAVING CYSTIC FIBROSIS USING IVACAFTOR AND LUMACAFTOR

U-1912 METHOD OF TREATING CYSTIC FIBROSIS IN A PATIENT, THE PATIENT HAVING THE F508DEL MUTATION IN CFTR, USING A DOSAGE UNIT AS DEFINED IN CLAIM 1 OF U.S. PATENT NO. 9,192,606

U-1913 TREATMENT OF PEDIATRIC PATIENTS WITH BILATERAL OTITIS MEDIA WITH EFFUSION UNDERGOING TYMPANOSTOMY TUBE PLACEMENT

U-1914 IN COMBINATION WITH RITUXIMAB, FOR THE TREATMENT OF PATIENTS WITH RELAPSED CHRONIC LYMPHOCYTIC LEUKEMIA (CLL)

U-1915 METHOD OF TREATING TYPE 2 DIABETES MELLITUS IN PATIENTS WITH SEVERE CHRONIC RENAL IMPAIRMENT AND WHO ARE INELIGIBLE FOR METFORMIN THERAPY BY ADMINISTERING LINAGLIPTIN

U-1916 PREVENTION OF NAUSEA AND VOMITING ASSOCIATED WITH CHEMOTHERAPY (CINV)

U-1917 TREATMENT OF EXOCRINE PANCREATIC CANCER THAT HAS PROGRESSED ON GEMCITABINE-BASED THERAPY, IN COMBINATION WITH 5-FLUOROURACIL AND LEUCOVORIN

U-1918 TREATMENT OF DUCHENNE MUSCULAR DYSTROPHY IN PATIENTS HAVING A MUTATION OF THE DMD GENE THAT IS AMENABLE TO EXON 51 SKIPPING

U-1919 RESTORING AN MRNA READING FRAME TO INDUCE DYSTROPHIN PROTEIN PRODUCTION IN PATIENTS HAVING A MUTATION OF THE DMD GENE THAT IS AMENABLE TO EXON 51 SKIPPING

U-1920 USE OF EXTENDED RELEASE ORAL 25-HYDROXYVITAMIN D3 IN TREATING SECONDARY HYPERPARATHYROIDISM IN ADULT PATIENTS HAVING CHRONIC KIDNEY DISEASE STAGE 3 OR STAGE 4

U-1921 MANAGEMENT OF PAIN SEVERE ENOUGH TO REQUIRE DAILY, AROUND-THE-CLOCK, LONG-TERM OPIOID TREATMENT AND FOR WHICH ALTERNATIVE TREATMENT OPTIONS ARE INADEQUATE BY PROVIDING AN ABUSE-DETERRENT ORAL CONTROLLED RELEASE COMBINATION DRUG PRODUCT

PATENT AND EXCLUSIVITY TERMS

ADB 135 of 225

PATENT USE

U-1922 INTRAVAGINAL PRASTERONE (DEHYDROEPIANDROSTERONE) AT A DAILY DOSE OF 6.5MG FOR THE TREATMENT OF DYSPAREUNIA, A SYMPTOM OF VULVAR AND VAGINAL ATROPHY, DUE TO MENOPAUSE

U-1923 IMPROVEMENT IN GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS INADEQUATELY CONTROLLED BY BASAL INSULIN OR LIXISENATIDE BY USE OF A PEN INJECTOR WITH A THREADED DRIVE SLEEVE

U-1924 KYPROLIS IS INDICATED IN COMBINATION WITH LENALIDOMIDE PLUS DEXAMETHASONE FOR THE TREATMENT OF PATIENTS WITH RELAPSED OR REFRACTORY MULTIPLE MYELOMA WHO HAVE RECEIVED ONE TO THREE LINES OF THERAPY

U-1925 USE OF AN AUTO INJECTOR TO ADMINISTER NALOXONE HCL

U-1926 METHOD OF TREATING, REDUCING THE INCIDENCE OF, OR PREVENTING AN ISCHEMIC EVENT IN A PATIENT UNDERGOING PCI BY ADMINISTERING INTRAVENOUSLY 30 UG/KG BOLUS BEFORE PCI AND CONTINUOUS INFUSION OF 4 UG/KG/MIN FOR AT LEAST 2 HOURS OR THE DURATION OF THE PCI

U-1927 TREATMENT OF PATIENTS WITH ANAPLASTIC LYMPHOMA KINASE (ALK)-POSITIVE METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) WHO HAVE PROGRESSED ON OR ARE INTOLERANT TO CRIZOTINIB

U-1928 RUBRACA IS INDICATED AS MONOTHERAPY FOR THE TREATMENT OF PATIENTS WITH DELETERIOUS BRCA MUTATION (GERMLINE AND/OR SOMATIC) ASSOCIATED ADVANCED OVARIAN CANCER WHO HAVE BEEN TREATED WITH TWO OR MORE CHEMOTHERAPIES.

U-1929 TREATMENT OF DIABETES MELLITUS WITH AN INHALED INSULIN TO IMPROVE GLYCEMIC CONTROL USING A DRY POWDER INHALATION SYSTEM COMPRISING AN INHALER, A CARTRIDGE AND A DRY POWDER MEDICAMENT COMPRISING INSULIN IN A SINGLE INHALATION

U-1930 METHOD OF AEROSOLIZING/DEAGGLOMERATING AN INSULIN DRY POWDER FOR USE IN TREATING DIABETES MELLITUS VIA ORAL INHALATION USING AN INHALER WITH A CARTRIDGE CONTAINING THE INSULIN DRY POWDER.

U-1931 PROPHYLAXIS OR TREATMENT OF VENOUS AND ARTERIAL THROMBOTIC DISEASE

U-1932 METHOD OF TREATING MILD TO MODERATE ATOPIC DERMATITIS.

U-1933 TREATMENT OF POSTOPERATIVE INFLAMMATION AND REDUCTION OF OCULAR PAIN IN PATIENTS WHO HAVE UNDERGONE CATARACT SURGERY

U-1934 TREATMENT OF NON-24-HOUR SLEEP-WAKE DISORDER BY AVOIDING THE USE OF TASIMELTEON IN COMBINATION WITH A STRONG CYP1A2 INHIBITOR

U-1935 REDUCTION OF THE RATE OF CARDIOVASCULAR DEATH, MYOCARDIAL INFARCTION, AND STROKE IN PATIENTS WITH A HISTORY OF MYOCARDIAL INFARCTION

U-1936 TREATMENT OF MYOCARDIAL INFARCTION AND STROKE IN PATIENTS WITH ACUTE CORONARY SYNDROME OR A HISTORY OF MYOCARDIAL INFARCTION

U-1937 TREATMENT OF MYOCARDIAL INFARCTION IN PATIENTS WITH ACUTE CORONARY SYNDROME OR A HISTORY OF MYOCARDIAL INFARCTION

U-1938 TREATMENT OF STROKE IN PATIENTS WITH ACUTE CORONARY SYNDROME OR A HISTORY OF MYOCARDIAL INFARCTION

U-1939 ADMINISTRATION ONCE DAILY WITHIN TWO HOURS AFTER WAKING IN THE MORNING FOR IMPROVEMENT OF GLYCEMIC CONTROL IN A TYPE 2 DIABETES PATIENT

U-1940 IMPROVEMENT IN THE APPEARANCE OF MODERATE TO SEVERE CONVEXITY OR FULLNESS ASSOCIATED WITH SUBMENTAL FAT IN ADULTS BY MEANS OF REDUCING SUBMENTAL FAT VOLUME AS DESCRIBED IN THE APPROVED LABELING

U-1941 TREATMENT OF INFANTILE-ONSET SPINAL MUSCULAR ATROPHY

U-1942 TREATMENT OF SPINAL MUSCULAR ATROPHY BY INCREASING EXON-7 INCLUSION IN SMN2 MRNA

U-1943 TREATMENT OF SPINAL MUSCULAR ATROPHY

U-1944 TREATMENT OF SPINAL MUSCULAR ATROPHY BY INHIBITING AN SMN2 PRE-MRNA INTRONIC SPLICING SILENCER SITE

U-1945 IMPROVEMENT OF GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS IN COMBINATION WITH METFORMIN.

U-1946 TREATMENT OF SMALL LYMPHOCYTIC LYMPHOMA

U-1947 TREATMENT OF MARGINAL ZONE LYMPHOMA

U-1948 A METHOD FOR TREATING CHRONIC MYELOID LEUKEMIA

U-1949 FOR USE IN THE TREATMENT OF PATIENTS WITH CHRONIC LYMPHOCYTIC LEUKEMIA (CLL)

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

- U-1950 TREATMENT OF PATIENTS WITH ADVANCED (METASTATIC) NON-SMALL CELL LUNG CANCER WHOSE DISEASE PROGRESSED DURING OR AFTER PLATINUM-BASED CHEMOTHERAPY
- U-1951 TREATMENT OF ONYCHOMYCOSIS OF A TOENAIL
- U-1952 FOR USE IN THE TREATMENT OF PATIENTS WITH INDOLENT B-CELL NON-HODGKIN LYMPHOMA
- U-1953 REDUCE THE RISK OF STROKE IN PATIENTS WITH NONVALVULAR ATRIAL FIBRILLATION WITH ONCE DAILY, RAPID-RELEASE TABLET ADMINISTERED FOR AT LEAST FIVE CONSECUTIVE DAYS
- U-1954 TREATMENT OF DEEP VEIN THROMBOSIS WITH ONCE DAILY, RAPID-RELEASE TABLET ADMINISTERED FOR AT LEAST FIVE CONSECUTIVE DAYS
- U-1955 TREATMENT OF PULMONARY EMBOLISM WITH ONCE DAILY, RAPID-RELEASE TABLET ADMINISTERED FOR AT LEAST FIVE CONSECUTIVE DAYS
- U-1956 FOLLOWING INITIAL 6 MONTHS TREATMENT FOR DEEP VEIN THROMBOSIS (DVT) AND/OR PULMONARY EMBOLISM (PE), REDUCTION IN THE RISK OF RECURRENCE OF DVT AND OF PE WITH ONCE DAILY, RAPID-RELEASE TABLET ADMINISTERED FOR AT LEAST FIVE CONSECUTIVE DAYS
- U-1957 PROPHYLAXIS OF DEEP VEIN THROMBOSIS, WHICH MAY LEAD TO PULMONARY EMBOLISM IN PATIENTS UNDERGOING KNEE OR HIP REPLACEMENT SURGERY, WITH ONCE DAILY, RAPID-RELEASE TABLET ADMINISTERED FOR AT LEAST FIVE CONSECUTIVE DAYS
- U-1958 FOR THE TREATMENT OF GENOTYPE 1, 2, 3 OR 4 CHRONIC HEPATITIS C VIRUS (HCV) INFECTION AS A COMPONENT OF A COMBINATION ANTIVIRAL TREATMENT REGIMEN WITH RIBAVIRIN
- U-1959 TOPICAL TREATMENT OF PERSISTENT FACIAL ERYTHEMA ASSOCIATED WITH ROSACEA IN ADULTS WITH 1% OXYMETAZOLINE HYDROCHLORIDE CREAM
- U-1960 USE OF THE ATYPICAL ANTIPSYCHOTIC ASENAPINE FOR TREATMENT OF SCHIZOPHRENIA IN ADULTS
- U-1961 USE OF THE ATYPICAL ANTIPSYCHOTIC ASENAPINE FOR TREATMENT OF MANIC OR MIXED EPISODES OF BIPOLAR I DISORDER: ACUTE MONOTHERAPY OF MANIC OR MIXED EPISODES (AGES 10 TO ADULT)
- U-1962 USE OF THE ATYPICAL ANTIPSYCHOTIC ASENAPINE FOR TREATMENT OF MANIC OR MIXED EPISODES OF BIPOLAR I DISORDER: MAINTENANCE MONOTHERAPY IN ADULTS
- U-1963 USE OF THE ATYPICAL ANTIPSYCHOTIC ASENAPINE FOR TREATMENT OF MANIC OR MIXED EPISODES OF BIPOLAR I DISORDER: AS ADJUNCTIVE TREATMENT TO LITHIUM OR VALPROATE IN ADULTS
- U-1964 ELEVATION OF INTRACELLULAR CGMP RESULTING IN INCREASED INTESTINAL FLUID AND ACCELERATED TRANSIT
- U-1965 FOR THE TREATMENT OF PULMONARY ARTERIAL HYPERTENSION (PAH) IN COMBINATION WITH TADALAFIL, WHEREIN THE WEIGHT RATIO OF AMBRISENTAN TO TADALAFIL IS ABOUT 1:2 TO ABOUT 1:3
- U-1966 USE OF THE ATYPICAL ANTIPSYCHOTIC ASENAPINE FOR TREATMENT OF MANIC OR MIXED EPISODES OF BIPOLAR I DISORDER: ACUTE MONOTHERAPY OF MANIC OR MIXED EPISODES IN PEDIATRIC PATIENTS AGE 10-17
- U-1967 METHOD OF TREATING TYPE 2 DIABETES IN PATIENTS WITH INSUFFICIENT GLYCEMIC CONTROL DESPITE THERAPY WITH ONE OR MORE CONVENTIONAL ANTIHYPERGLYCEMIC AGENTS BY ADMINISTERING LINAGLIPTIN IN COMBINATION WITH METFORMIN
- U-1968 METHOD OF TREATING TYPE 2 DIABETES IN PATIENTS WHO HAVE NOT BEEN PREVIOUSLY TREATED WITH AN ANTIHYPERGLYCEMIC AGENT BY ADMINISTERING LINAGLIPTIN IN COMBINATION WITH METFORMIN
- U-1969 TOPICAL TREATMENT OF ONYCHOMYCOSIS OF THE TOENAIL(S) DUE TO TRICHOPHYTON RUBRUM AND TRICHOPHYTON MENTAGROPHYTES
- U-1970 TREATMENT OF ONYCHOMYCOSIS OF A TOENAIL CAUSED BY TRICHOPHYTON RUBRUM OR TRICHOPHYTON MENTAGROPHYTES
- U-1971 FOR THE TREATMENT OF PATIENTS WITH CHRONIC LYMPHOCYTIC LEUKEMIA
- U-1972 FOR THE TREATMENT OF PATIENTS WITH INDOLENT B-CELL NON-HODGKIN LYMPHOMA
- U-1973 METHOD OF TREATING CYSTIC FIBROSIS USING N-(5-HYDROXY-2,4-DITERT-BUTYL-PHENYL)-4-OXO-1H-QUINOLINE-3-CARBOXAMIDE AND 3-(6-(1-2,2-DIFLUOROBENZO[D][1,3]DIOXOL-5-YL) CYCLOPROPANECARBOXAMIDO)-3-METHYLPYRIDIN-2-YL)BENZOIC ACID
- U-1974 TREATMENT OF HALLUCINATIONS AND DELUSIONS ASSOCIATED WITH PARKINSON'S DISEASE PSYCHOSIS
- U-1975 METHOD OF INCREASING EYELASH GROWTH WITH BIMATOPROST

PATENT AND EXCLUSIVITY TERMS

ADB 137 of 225

PATENT USE

U-1976 METHOD FOR TREATING TYPE 2 DIABETES MELLITUS (T2DM) IN PATIENTS WHO HAVE INADEQUATE CONTROL WITH DAPAGLIFLOZIN

U-1977 METHOD FOR TREATING TYPE 2 DIABETES MELLITUS (T2DM) IN PATIENTS WHO ARE ALREADY TREATED WITH DAPAGLIFLOZIN AND SAXAGLIPTIN

U-1978 TREATMENT OF ADVANCED PROSTATE CANCER WITH A REDUCED LIKELIHOOD OF CAUSING A GONADOTROPHIN RELEASING HORMONE AGONIST SIDE-EFFECT

U-1979 THE TREATMENT OF CARCINOID SYNDROME DIARRHEA IN COMBINATION WITH SOMATOSTATIN ANALOG (SSA) THERAPY IN ADULTS INADEQUATELY CONTROLLED BY SSA THERAPY

U-1980 A METHOD OF TREATING NOCTURIA DUE TO NOCTURNAL POLYURIA IN ADULTS

U-1981 IN COMBINATION WITH AN AROMATASE INHIBITOR AS INITIAL ENDOCRINE-BASED THERAPY FOR TREATMENT OF POSTMENOPAUSAL WOMEN WITH HORMONE RECEPTOR (HR)-POSITIVE, HUMAN EPIDERMAL GROWTH FACTOR RECEPTOR 2 (HER-2)-NEGATIVE ADVANCED OR METASTATIC BREAST CANCER

U-1982 USE OF REVLIMID (LENALIDOMIDE) FOR TREATMENT OF PATIENTS WITH TRANSFUSION-DEPENDENT ANEMIA DUE TO LOW-OR INTERMEDIATE-1-RISK MYELODYSPLASTIC SYNDROMES ASSOCIATED WITH A DELETION 5Q ABNORMALITY WITH OR WITHOUT ADDITIONAL CYTOGENETIC ABNORMALITIES

U-1983 USE OF REVLIMID (LENALIDOMIDE) FOR THE TREATMENT OF PATIENTS WITH MANTLE CELL LYMPHOMA WHOSE DISEASE HAS RELAPSED OR PROGRESSED AFTER TWO PRIOR THERAPIES, ONE OF WHICH INCLUDED BORTEZOMIB

U-1984 USE OF REVLIMID (LENALIDOMIDE) FOR THE TREATMENT OF PATIENTS WITH MULTIPLE MYELOMA, IN COMBINATION WITH DEXAMETHASONE

U-1985 USE OF REVLIMID (LENALIDOMIDE) FOR THE TREATMENT OF PATIENTS WITH MULTIPLE MYELOMA, AS MAINTENANCE FOLLOWING AUTOLOGOUS HEMATOPOIETIC STEM CELL TRANSPLANTATION (AUTO-HSCT)

U-1986 USE OF REVLIMID (LENALIDOMIDE) FOR THE TREATMENT OF PATIENTS WITH MULTIPLE MYELOMA, IN COMBINATION WITH DEXAMETHASONE, WHEREIN THOSE PATIENTS HAVE NOT RECEIVED PREVIOUS TREATMENT FOR MULTIPLE MYELOMA

U-1987 METHOD OF CONTROLLING GLYCEMIA IN DIABETICS BY ADMINISTERING AN INITIAL DOSE OF INSULIN-FDKP WITH A MEAL; DETERMINING BLOOD GLUCOSE LEVEL 1-2 HRS AFTER AND ADMINISTERING A SUPPLEMENTAL DOSE OF INSULIN-FDKP IF POSTPRANDIAL GLUCOSE LEVEL IS >140 MG/DL

U-1988 METHOD TO TREAT INFANTILE HEMANGIOMA

U-1989 INTRAVITREAL TREATMENT OF MACULAR EDEMA FOLLOWING BRANCH RETINAL VEIN OCCLUSION (BRVO) OR CENTRAL RETINAL VEIN OCCLUSION (CRVO)

U-1990 INTRAVITREAL TREATMENT OF DIABETIC MACULAR EDEMA

U-1991 REDUCTION OF MORTALITY IN ACUTE MYOCARDIAL INFARCTION

U-1992 USE OF TROKENDI XR FOR PROPHYLACTIC TREATMENT OF MIGRAINE

U-1993 ADJUNCTIVE TREATMENT TO LEVODOPA/CARBIDOPA IN PATIENTS WITH PARKINSON'S DISEASE EXPERIENCING "OFF" EPISODES

U-1994 REDUCTION IN RISK OF OVERT HEPATIC ENCEPHALOPATHY (HE) IN ADULTS

U-1995 TREATMENT OF TARDIVE DYSKINESIA

U-1996 IMPROVEMENT OF GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS

U-1997 IMPROVEMENT OF GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS IN COMBINATION WITH METFORMIN AND/OR A PPAR-GAMMA AGONIST AND/OR SULFONYLUREA AND/OR INSULIN

U-1998 TREATING HR-POS., HER2-NEG. ADVANCED OR METASTATIC BREAST CANCER WITH PALBOCICLIB IN COMBO WITH AN AROMATASE INHIBITOR AS INITIAL ENDOCRINE BASED THERAPY IN POSTMENOPAUSAL WOMEN OR FULVESTRANT IN WOMEN WITH DISEASE PROGRESSION AFTER ENDOCRINE THERAPY

U-1999 CHRONIC IDIOPATHIC CONSTIPATION

U-2000 MANAGEMENT OF MODERATE TO SEVERE PAIN AS AN ADJUNCT TO OPIOID ANALGESICS

U-2001 USE FOR THE TREATMENT OF ASTHMA IN PATIENTS 6 YEARS OF AGE AND OLDER

U-2002 USE FOR MAINTENANCE TREATMENT OF CHRONIC OBSTRUCTIVE PULMONARY DISEASE

U-2003 A METHOD OF POSITIONING AN INTRAUTERINE SYSTEM BY HOLDING AN INSERTER HANDLE WITH ONE HAND, ADVANCING THE INSERTER THROUGH THE CERVIX AND INTO THE UTERUS, AND RETRACTING A SLIDER ON THE HANDLE TO RELEASE THE INTRAUTERINE SYSTEM

PATENT AND EXCLUSIVITY TERMS

ADB 138 of 225

PATENT USE

U-2004 REPLACEMENT THERAPY FOR ORAL CARBAMAZEPINE IN ADULTS WITH PARTIAL SEIZURES WITH COMPLEX SYMPTOMATOLOGY

U-2005 REPLACEMENT THERAPY FOR ORAL CARBAMAZEPINE IN ADULTS WITH GENERALIZED TONIC-CLONIC SEIZURES

U-2006 REPLACEMENT THERAPY FOR ORAL CARBAMAZEPINE IN ADULTS WITH MIXED SEIZURE PATTERNS THAT INCLUDE PARTIAL SEIZURES WITH COMPLEX SYMPTOMATOLOGY, GENERALIZED TONIC-CLONIC SEIZURES, OR OTHER PARTIAL OR GENERALIZED SEIZURES

U-2007 TREATMENT OF ADULT PATIENTS WITH NEWLY DIAGNOSED ACUTE MYELOID LEUKEMIA (AML) WHO ARE FLT3 MUTATION-POSITIVE, IN COMBINATION WITH STANDARD CYTARABINE AND DAUNORUBICIN INDUCTION AND CYTARABINE CONSOLIDATION CHEMOTHERAPY

U-2008 TREATMENT OF ADULT PATIENTS WITH AGGRESSIVE SYSTEMIC MASTOCYTOSIS (ASM), SYSTEMIC MASTOCYTOSIS WITH ASSOCIATED HEMATOLOGICAL NEOPLASM (SM-AHN), OR MAST CELL LEUKEMIA (MCL)

U-2009 METHOD OF TREATING POSTMENOPAUSAL WOMEN WITH OSTEOPOROSIS AT HIGH RISK FOR FRACTURE.

U-2010 ACUTE TREATMENT OF MIGRAINE BY DELIVERING A POWDERED SUBSTANCE COMPRISING SUMATRIPTAN VIA A BREATH-POWERED DELIVERY DEVICE

U-2011 TREATMENT OF MIGRAINE VIA DELIVERY OF SUMATRIPTAN VIA THE NASAL CAVITY

U-2012 A METHOD FOR TREATING OVARIAN CANCER BY ADMINISTERING RUCAPARIB, WHEREIN THE CANCER IS ASSOCIATED WITH A DELETERIOUS BRCA MUTATION

U-2013 TREATMENT OF AMYOTROPHIC LATERAL SCLEROSIS (ALS)

U-2014 A METHOD OF TREATING SECONDARY HYPERPARATHYROIDISM (SHPT)

U-2015 SODIUM THIOSULFATE INJECTION IS INDICATED FOR SEQUENTIAL USE WITH SODIUM NITRITE FOR THE TREATMENT OF ACUTE CYANIDE POISONING

U-2016 TREATMENT FOR ONYCHOMYCOSIS THAT IS TINEA UNGUIUM

U-2017 TREATMENT OF OPIOID DEPENDENCE

U-2018 MANAGEMENT OF MILD TO MODERATE PAIN, MANAGEMENT OF MODERATE TO SEVERE PAIN AS AN ADJUNCT TO OPIOID ANALGESICS IN A CRITICALLY ILL PATIENT WITH INTRAVENOUS IBUPROFEN IN NEED THEREOF

U-2019 METHOD OF DELIVERING TO A PATIENT WITH DIABETES MELLITUS IN A SINGLE INHALATION, GREATER THAN 75% OF A DRY POWDER DOSE COMPRISING INSULIN AND FUMARYL DIKETOPIPERAZINE USING A HIGH RESISTANCE TO FLOW DRY POWDER INHALER.

U-2020 MEKINIST IS INDICATED, IN COMBINATION WITH DABRAFENIB, FOR THE TREATMENT OF PATIENTS WITH METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) WITH BRAF V600E MUTATION AS DETECTED BY AN FDA-APPROVED TEST

U-2021 METHOD OF ADMINISTERING LEVETIRACETAM UNDER FASTED CONDITIONS

U-2022 METHOD OF ADMINISTERING LEVETIRACETAM UNDER FED CONDITIONS

U-2023 A METHOD OF INCREASING THE BIOAVAILABILITY OF GUAIFENESIN IN A SOLUTION CONTAINING 54% TO 66% BY WEIGHT OF PROPYLENE GLYCOL AND GLYCEROL, WHEREIN THE METHOD INCREASES THE CMAX BY AT LEAST 1.5 AND/OR INCREASES THE AUC (0-INF) BY AT LEAST 1.4

U-2024 METHOD FOR TRANSDERMALLY DELIVERING A DRUG TO A USER IN NEED THEREOF

U-2025 TREATMENT OF ATTENTION DEFICIT HYPERACTIVITY DISORDER

U-2026 TAFINLAR(R) IS INDICATED, IN COMBINATION WITH TRAMETINIB, FOR THE TREATMENT OF PATIENTS WITH UNRESECTABLE OR METASTATIC MELANOMA WITH BRAF V600E OR V600K MUTATIONS, AS DETECTED BY AN FDA-APPROVED TEST.

U-2027 TAFINLAR(R) IS INDICATED, IN COMBINATION WITH TRAMETINIB, FOR THE TREATMENT OF PATIENTS WITH METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) WITH BRAF V600E MUTATION AS DETECTED BY AN FDA-APPROVED TEST.

U-2028 TREATMENT OF ACUTE BACTERIAL SKIN AND SKIN STRUCTURE INFECTIONS CAUSED BY DESIGNATED SUSCEPTIBLE BACTERIA IN ADULTS

U-2029 PREVENTING CONDITION CHARACTERIZED BY UNDESIRED THROMBOSIS

U-2030 PROPHYLAXIS OF VENOUS THROMBOSIS

U-2031 TAFINLAR IS INDICATED AS A SINGLE AGENT FOR THE TREATMENT OF PATIENTS WITH UNRESECTABLE OR METASTATIC MELANOMA WITH BRAF V600E MUTATION AS DETECTED BY AN FDA-APPROVED TEST

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

- U-2032 TAFINLAR IS INDICATED, IN COMBINATION WITH TRAMETINIB, FOR THE TREATMENT OF PATIENTS WITH UNRESECTABLE OR METASTATIC MELANOMA WITH BRAF V600E OR V600K MUTATIONS, AS DETECTED BY AN FDA-APPROVED TEST.
- U-2033 MEKINIST IS INDICATED, AS A SINGLE AGENT OR IN COMBINATION WITH DABRAFENIB, FOR THE TREATMENT OF PATIENTS WITH UNRESECTABLE OR METASTATIC MELANOMA WITH BRAF V600E OR V600K MUTATIONS
- U-2034 INHIBITING COAGULATION
- U-2035 PROPHYLAXIS OF VENOUS THROMBOEMBOLISM
- U-2036 A METHOD OF TREATING PULMONARY HYPERTENSION COMPRISING PARENTERALLY ADMINISTERING A FORMULATION COMPRISING A) 0.1 TO 5% W/V OF TREPROSTINIL OR A PHARMACEUTICALLY ACCEPTABLE SALT THEREOF AND B) A CITRATE BUFFER
- U-2037 MEKINIST IS INDICATED, AS A SINGLE AGENT OR IN COMBINATION WITH DABRAFENIB, FOR THE TREATMENT OF PATIENTS WITH UNRESECTABLE OR METASTATIC MELANOMA WITH BRAF V600E OR V600K MUTATIONS, AS DETECTED BY AN FDA-APPROVED TEST
- U-2038 MEKINIST IS INDICATED, IN COMBINATION WITH DABRAFENIB, FOR THE TREATMENT OF PATIENTS WITH METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) WITH BRAF V600E MUTATION AS DETECTED BY AN FDA-APPROVED TEST
- U-2039 TREATMENT OF ADULT PATIENTS WITH CHRONIC HCV INFECTION WHO HAVE GENOTYPE 1, 2, 3, 4, 5, OR 6 INFECTION AND HAVE PREVIOUSLY BEEN TREATED WITH AN HCV REGIMEN CONTAINING AN NS5A INHIBITOR
- U-2040 TREATMENT OF ADULT PATIENTS WITH CHRONIC HCV INFECTION WHO HAVE GENOTYPE 1A OR 3 INFECTION AND HAVE PREVIOUSLY BEEN TREATED WITH AN HCV REGIMEN CONTAINING SOFOSBUVIR WITHOUT AN NS5A INHIBITOR
- U-2041 TREATMENT OF PARTIAL-ONSET SEIZURES
- U-2042 DISCONTINUING ADMINISTRATION OF FLUVOXAMINE TO AVOID DRUG INTERACTIONS WITH PIRFENIDONE AND THEN ADMINISTERING PIRFENIDONE
- U-2043 EXTENDED ADJUVANT TREATMENT OF ADULT PATIENTS WITH EARLY STAGE HER2-OVEREXPRESSED/AMPLIFIED BREAST CANCER, TO FOLLOW ADJUVANT TRASTUZUMAB BASE THERAPY
- U-2044 DOSE REDUCTION OF PIRFENIDONE BY ABOUT ONE HALF DURING CONCURRENT ADMINISTRATION OF CIPROFLOXACIN AT A DOSE OF 750 MG TWICE DAILY (1500 MG/DAY) TO REDUCE DRUG INTERACTIONS IN TREATMENT OF A FIBROTIC, INFLAMMATORY, OR AUTOIMMUNE DISORDER
- U-2045 ADMINISTRATION OF PIRFENIDONE AND AVOIDING CONCURRENT ADMINISTRATION OF CIPROFLOXACIN AT A DOSE OF 750 MG TO REDUCE DRUG INTERACTIONS IN TREATMENT OF A FIBROTIC, INFLAMMATORY, OR AUTOIMMUNE DISORDER
- U-2046 ADMINISTERING PIRFENIDONE WHILE AVOIDING CONCOMITANT USE OF A CYP1A2 INHIBITOR THAT IS A MODERATE TO STRONG INHIBITOR OF BOTH CYP1A2 AND ANOTHER CYP ENZYME SELECTED FROM CYP2C9, CYP2C19, AND CYP2D6
- U-2047 ADMINISTERING PIRFENIDONE CONCURRENTLY WITH FLUVOXAMINE, THE PIRFENIDONE AT A DOSE OF ABOUT 801 MG/DAY TO REDUCE DRUG INTERACTIONS WITH FLUVOXAMINE
- U-2048 ADMINISTERING PIRFENIDONE WHILE AVOIDING CO-ADMINISTRATION OF A STRONG CYP1A2 INHIBITOR TO AVOID DRUG INTERACTIONS WITH PIRFENIDONE
- U-2049 DISCONTINUING ADMINISTRATION OF A STRONG CYP1A2 INHIBITOR TO AVOID DRUG INTERACTIONS WITH PIRFENIDONE AND THEN ADMINISTERING PIRFENIDONE
- U-2050 ADMINISTERING PIRFENIDONE WHILE AVOIDING CO-ADMINISTRATION OF FLUVOXAMINE TO AVOID DRUG INTERACTIONS WITH PIRFENIDONE
- U-2051 DISCONTINUING SMOKING TO AVOID REDUCED PIRFENIDONE EFFICACY AND THEN ADMINISTERING PIRFENIDONE
- U-2052 DISCONTINUING ADMINISTRATION OF A STRONG CYP1A2 INDUCER TO AVOID REDUCED PIRFENIDONE EFFICACY AND THEN ADMINISTERING PIRFENIDONE
- U-2053 ADMINISTERING PIRFENIDONE WHILE AVOIDING CONCOMITANT ADMINISTRATION OF A STRONG INDUCER OF CYP1A2, INCLUDING CIGARETTE SMOKE, TO AVOID REDUCED PIRFENIDONE EFFICACY
- U-2054 ADMINISTERING PIRFENIDONE WHILE AVOIDING CONCOMITANT ADMINISTRATION OF A STRONG INDUCER OF CYP1A2 TO AVOID REDUCED PIRFENIDONE EFFICACY
- U-2055 DOSING OF AT LEAST 1600 MG/DAY FOLLOWING GRADE 2 LIVER ABNORMALITY IN BIOMARKER AST AND/OR ALT AFTER PIRFENIDONE ADMINISTRATION IN TREATMENT OF IPF
- U-2056 DOSAGE MODIFICATION FOLLOWING GRADE 2 ABNORMALITY IN BIOMARKER AST AND/OR ALT AFTER PIRFENIDONE ADMINISTRATION, BY ADMINISTERING SUB-1600 MG/DAY, FOLLOWING BY

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

- ADMINISTERING AT LEAST 1600 MG/DAY IN TREATMENT OF IPF
- U-2057 DOSING 2403 MG/DAY PIRFENIDONE FOLLOWING GRADE 2 ABNORMALITY IN BIOMARKER AST AND/OR ALT AFTER PIRFENIDONE ADMINISTRATION IN TREATMENT OF IPF
- U-2058 DOSAGE MODIFICATION FOLLOWING GRADE 2 ABNORMALITY IN BIOMARKER AST AND/OR ALT AFTER PIRFENIDONE ADMINISTRATION, BY ADMINISTERING SUB-2400MG/DAY DOSE, FOLLOWED BY ADMINISTERING 2403MG/DAY IN TREATMENT OF IPF
- U-2059 DOSAGE MODIFICATION FOLLOWING GRADE 2 ABNORMALITY IN BIOMARKER AST AND/OR ALT AFTER PIRFENIDONE ADMINISTRATION, BY DISCONTINUING PIRFENIDONE UNTIL BIOMARKERS OF LIVER FUNCTION ARE WITHIN NORMAL LIMITS, FOLLOWED BY FULL DAILY DOSE IN TREATMENT OF IPF
- U-2060 DOSAGE MODIFICATION FOLLOWING GRADE 2 ABNORMALITY IN BIOMARKER AST AND/OR ALT AFTER PIRFENIDONE ADMINISTRATION, BY DISCONTINUING PIRFENIDONE UNTIL BIOMARKERS OF LIVER FUNCTION ARE WITHIN NORMAL LIMITS, THEN AT LEAST 1600MG/DAY IN TREATMENT OF IPF
- U-2061 DOSING OF AT LEAST 1600 MG/DAY FOLLOWING GRADE 2 ABNORMALITY IN LIVER FUNCTION BIOMARKER AFTER PIRFENIDONE ADMINISTRATION IN TREATMENT OF IPF
- U-2062 DOSAGE MODIFICATION FOLLOWING GRADE 2 ABNORMALITY IN LIVER FUNCTION BIOMARKER AFTER PIRFENIDONE ADMINISTRATION, BY ADMINISTERING SUB-1600 MG/DAY DOSE, FOLLOWED BY ADMINISTERING AT LEAST 1600 MG/DAY DOSE IN TREATMENT OF IPF
- U-2063 DOSAGE MODIFICATION FOLLOWING GRADE 2 ABNORMALITY IN LIVER FUNCTION BIOMARKER AFTER PIRFENIDONE ADMINISTRATION, BY DISCONTINUING PIRFENIDONE UNTIL BIOMARKERS ARE WITHIN NORMAL LIMITS, FOLLOWED BY ADMINISTERING AT LEAST 1600 MG/DAY IN TREATMENT OF IPF
- U-2064 DOSING AT LEAST 1602 MG/DAY FOLLOWING GRADE 2 ABNORMALITY IN LIVER FUNCTION BIOMARKER ALT OR AST AFTER PIRFENIDONE ADMINISTRATION
- U-2065 FULL DAILY DOSING FOLLOWING GRADE 2 ABNORMALITY IN LIVER FUNCTION BIOMARKER ALT OR AST AFTER PIRFENIDONE ADMINISTRATION
- U-2066 DOSAGE MODIFICATION FOLLOWING GRADE 2 ABNORMALITY IN LIVER FUNCTION BIOMARKER ALT OR AST AFTER PIRFENIDONE ADMINISTRATION, BY ADMINISTERING SUB-2400 MG/DAY DOSE, FOLLOWED BY FULL DAILY DOSE
- U-2067 DOSAGE MODIFICATION FOLLOWING GRADE 2 ABNORMALITY IN LIVER FUNCTION BIOMARKER ALT OR AST AFTER PIRFENIDONE ADMINISTRATION, BY DISCONTINUING PIRFENIDONE, FOLLOWED BY ADMINISTERING AT LEAST 1602 MG/DAY
- U-2068 DOSAGE MODIFICATION FOLLOWING GRADE 2 ABNORMALITY IN LIVER FUNCTION BIOMARKER ALT OR AST AFTER PIRFENIDONE ADMINISTRATION, BY DISCONTINUING PIRFENIDONE UNTIL BIOMARKERS OF LIVER FUNCTION ARE WITHIN NORMAL LIMITS, FOLLOWED BY FULL DAILY DOSE
- U-2069 DOSAGE MODIFICATION FOLLOWING GRADE 2 ABNORMALITY IN LIVER FUNCTION BIOMARKER ALT OR AST AFTER PIRFENIDONE ADMINISTRATION, BY ADMINISTERING A SUB-1600 MG/DAY DOSE, FOLLOWED BY ADMINISTERING AT LEAST 1602 MG/DAY
- U-2070 DOSAGE MODIFICATION FOLLOWING GRADE 2 ABNORMALITY IN BIOMARKER ALT OR AST AFTER PIRFENIDONE ADMINISTRATION, BY DISCONTINUING PIRFENIDONE UNTIL BIOMARKERS OF LIVER FUNCTION ARE WITHIN NORMAL LIMITS, THEN SUB-1600 MG/DAY, THEN AT LEAST 1602 MG/DAY
- U-2071 FULL DAILY DOSING FOLLOWING GRADE 2 ABNORMALITY IN LIVER FUNCTION BIOMARKER AFTER PIRFENIDONE ADMINISTRATION IN TREATMENT OF IPF
- U-2072 FULL DAILY DOSING FOLLOWING GRADE 2 ABNORMALITY IN LIVER FUNCTION BIOMARKER AFTER PIRFENIDONE ADMINISTRATION IN TREATMENT OF IPF
- U-2073 DOSAGE MODIFICATION FOLLOWING GRADE 2 ABNORMALITY IN LIVER FUNCTION BIOMARKER AFTER PIRFENIDONE ADMINISTRATION, BY DISCONTINUING PIRFENIDONE UNTIL BIOMARKERS ARE WITHIN NORMAL LIMITS, FOLLOWED BY ADMINISTERING FULL DAILY DOSE IN TREATMENT OF IPF
- U-2074 DOSING 1602 MG/DAY PIRFENIDONE FOLLOWING GRADE 2 ABNORMALITY IN LIVER FUNCTION BIOMARKER AFTER PIRFENIDONE ADMINISTRATION IN TREATMENT OF IPF
- U-2075 DOSAGE MODIFICATION FOLLOWING GRADE 2 ABNORMALITY IN LIVER FUNCTION BIOMARKER AFTER PIRFENIDONE ADMINISTRATION, BY DISCONTINUING PIRFENIDONE UNTIL BIOMARKERS ARE WITHIN NORMAL LIMITS FOLLOWED BY ADMINISTERING 1602 MG/DAY IN TREATMENT OF IPF
- U-2076 DOSAGE MODIFICATION FOLLOWING GRADE 2 ABNORMALITY IN LIVER FUNCTION BIOMARKER AFTER PIRFENIDONE ADMINISTRATION, BY ADMINISTERING 801 MG/DAY FOLLOWED BY ADMINISTERING 1602 MG/DAY IN TREATMENT OF IPF

PATENT AND EXCLUSIVITY TERMS

ADB 141 of 225

PATENT USE

- U-2077 DOSAGE MODIFICATION FOLLOWING GRADE 2 ABNORMALITY IN LIVER FUNCTION BIOMARKER AFTER PIRFENIDONE ADMINISTRATION, BY ADMINISTERING SUB-2400 MG/DAY DOSE THEN FULL DAY DAILY DOSE IN TREATMENT OF IPF
- U-2078 DOSAGE MODIFICATION FOLLOWING GRADE 2 ABNORMALITY IN LIVER FUNCTION BIOMARKER AFTER PIRFENIDONE ADMINISTRATION, BY DISCONTINUING PIRFENIDONE UNTIL BIOMARKERS ARE WITHIN LIMITS, THEN SUB-2400MG/DAY DOSE, THEN FULL DAILY DOSE IN TREATMENT OF IPF
- U-2079 PIRFENIDONE DOSE ESCALATION REGIMEN FOR TREATMENT OF FIBROSIS AS 801 MG/DAY FOR DAYS 1-7 OF THE REGIMEN, 1602 MG/DAY FOR DAYS 8-14 OF THE REGIMEN, AND 2403 MG/DAY FOR AT LEAST DAY 15 OF THE REGIMEN
- U-2080 PIRFENIDONE DOSE ESCALATION REGIMEN FOR TREATMENT OF IPF AS 801 MG/DAY FOR DAYS 1-7 OF THE REGIMEN, 1602 MG/DAY FOR DAYS 8-14 OF THE REGIMEN, AND 2403 MG/DAY FOR AT LEAST DAY 15 OF THE REGIMEN
- U-2081 DISCONTINUING USE OF A CYP1A2 INHIBITOR THAT IS A MODERATE TO STRONG INHIBITOR OF BOTH CYP1A2 AND ANOTHER CYP ENZYME SELECTED FROM CYP2C9, CYP2C19, AND CYP2D6 AND THEN ADMINISTERING PIRFENIDONE
- U-2082 MODIFYING PIRFENIDONE ADMINISTRATION FROM A DOSE OF ABOUT 2400 MG/DAY DOWNWARD BY ABOUT 1600 MG/DAY WHILE CO-ADMINISTERING FLUVOXAMINE TO REDUCE DRUG INTERACTIONS WITH FLUVOXAMINE
- U-2083 DOSAGE MODIFICATION FOLLOWING GRADE 2 ABNORMALITY IN LIVER FUNCTION BIOMARKER AFTER PIRFENIDONE ADMINISTRATION, BY DISCONTINUING PIRFENIDONE UNTIL BIOMARKERS ARE WITHIN NORMAL LIMITS, FOLLOWED BY 801 MG/DAY, DOSE, THEN 1602 MG/DAY IN TREATMENT OF IPF
- U-2084 TREATMENT OF SEVERE CHRONIC PAIN VIA INTRATHECAL INFUSION OF ZICONOTIDE IN PATIENTS ALSO RECEIVING MORPHINE
- U-2085 TREATMENT OF NON-24-HOUR SLEEP-WAKE DISORDER BY AVOIDING THE USE OF TASIMELTEON IN COMBINATION WITH RIFAMPIN
- U-2086 A METHOD FOR ADMINISTERING ESTRADIOL COMPRISING A MONOLITHIC TRANSDERMAL DRUG DELIVERY SYSTEM CONSISTING OF (I) A BACKING LAYER AND (II) A SINGLE ADHESIVE POLYMER MATRIX LAYER AS CLAIMED IN US PATENT NO. 9730900
- U-2087 TREATMENT OF RELAPSED OR REFRACTORY ACUTE MYELOID LEUKEMIA (AML) WITH AN ISOCITRATE DEHYDROGENASE-2 (IDH2) MUTATION
- U-2088 TREATMENT OF PARTIAL-ONSET SEIZURES WITH OR WITHOUT SECONDARILY GENERALIZED SEIZURES IN PATIENTS WITH EPILEPSY
- U-2089 TREATMENT OF PRIMARY GENERALIZED TONIC-CLONIC SEIZURES AS ADJUNCTIVE THERAPY IN PATIENTS WITH EPILEPSY
- U-2090 FOR THE TREATMENT OF ADULTS WITH NEWLY-DIAGNOSED THERAPY-RELATED ACUTE MYELOID LEUKEMIA (T-AML) OR AML WITH MYELODYSPLASIA-RELATED CHANGES (AML-MRC)
- U-2091 TREATMENT OF METASTATIC ADENOCARCINOMA OF THE PANCREAS THAT HAS PROGRESSED ON GEMCITABINE-BASED THERAPY, IN COMBINATION WITH 5-FLUOROURACIL AND LEUCOVORIN, IN A PATIENT NOT HOMOZYGOUS FOR THE UGT1A1*28 ALLELE
- U-2092 METHOD FOR CONFIRMING DOSE DELIVERY
- U-2093 TREATMENT OF TYPE II SPINAL MUSCULAR ATROPHY
- U-2094 TREATMENT OF TYPE III SPINAL MUSCULAR ATROPHY
- U-2095 MITOSOL IS AN ANTIMETABOLITE INDICATED AS AN ADJUNCT TO AB EXTERNO GLAUCOMA SURGERY. IT IS INTENDED FOR TOPICAL APPLICATION TO THE SITE OF GLAUCOMA FILTRATION SURGERY
- U-2096 SOTYLIZE IS INDICATED FOR THE MAINTENANCE OF NORMAL SINUS RHYTHM [DELAY IN TIME TO RECURRENCE OF ATRIAL FIBRILLATION/ATRIAL FLUTTER (AFIB/AFL)] IN PATIENTS WITH SYMPTOMATIC AFIB/AFL WHO ARE CURRENTLY IN SINUS RHYTHM
- U-2097 TREATMENT OF DMD IN PATIENTS HAVING A MUTATION OF THE DMD GENE THAT IS AMENABLE TO EXON 51 SKIPPING
- U-2098 INCREASING PRODUCTION OF FUNCTIONAL DYSTROPHIN PROTEIN IN DMD PATIENTS HAVING A MUTATION OF THE DMD GENE THAT IS AMENABLE TO EXON 51 SKIPPING
- U-2099 INDICATED FOR THE LONG-TERM, ONCE-DAILY, MAINTENANCE TREATMENT OF AIRFLOW OBSTRUCTION IN PATIENTS WITH CHRONIC OBSTRUCTIVE PULMONARY DISEASE (COPD), INCLUDING BRONCHITIS AND/OR EMPHYSEMA
- U-2100 INDICATED FOR THE ONCE-DAILY TREATMENT OF ASTHMA IN PATIENTS 18 YEARS AND OLDER

PATENT AND EXCLUSIVITY TERMS

ADB 142 of 225

PATENT USE

- U-2101 MAINTENANCE TREATMENT OF RECURRENT EPITHELIAL OVARIAN, FALLOPIAN TUBE OR PRIMARY PERITONEAL CANCER, WHO ARE IN A COMPLETE OR PARTIAL RESPONSE TO PLATINUM-BASED CHEMOTHERAPY
- U-2102 TREATMENT OF DELETERIOUS OR SUSPECTED DELETERIOUS GERMLINE BRCA-MUTATED ADVANCED OVARIAN CANCER WHO HAVE BEEN TREATED WITH THREE OR MORE PRIOR LINES OF CHEMOTHERAPY BASED ON AN FDA-APPROVED COMPANION DIAGNOSTIC FOR LYNPARZA
- U-2103 MAINTENANCE TREATMENT OF BRCA-MUTATED RECURRENT EPITHELIAL OVARIAN, FALLOPIAN TUBE OR PRIMARY PERITONEAL CANCER, WHO ARE IN A COMPLETE OR PARTIAL RESPONSE TO PLATINUM-BASED CHEMOTHERAPY
- U-2104 TREATMENT OF HYPERURICEMIA ASSOCIATED WITH GOUT IN PATIENTS WHO HAVE NOT ACHIEVED TARGET SERUM URIC ACID LEVELS WITH A MEDICALLY APPROPRIATE DAILY DOSE OF ALLOPURINOL ALONE
- U-2105 TREATMENT OF DYSKINESIA IN PATIENTS WITH PARKINSON'S DISEASE RECEIVING IMMEDIATE RELEASE LEVODOPA-BASED THERAPY, WITH OR WITHOUT CONCOMITANT DOPAMINERGIC MEDICATIONS
- U-2106 TREATMENT OF DYSKINESIA IN PATIENTS WITH PARKINSON'S DISEASE RECEIVING LEVODOPA-BASED THERAPY, WITH OR WITHOUT CONCOMITANT DOPAMINERGIC MEDICATIONS
- U-2107 TREATMENT OF LOCALLY RECURRENT OR METASTATIC, PROGRESSIVE, DIFFERENTIATED THYROID CARCINOMA REFRACTORY TO RADIOACTIVE IODINE TREATMENT
- U-2108 TREATMENT OF HORMONE RECEPTOR POSITIVE HER2-NEGATIVE ADVANCED BREAST CANCER IN POSTMENOPAUSAL WOMEN NOT PREVIOUSLY TREATED WITH ENDOCRINE THERAPY
- U-2109 CAROSPIR IS INDICATED FOR TREATMENT OF NYHA CLASS III-IV HEART FAILURE AND REDUCED EJECTION FRACTION TO INCREASE SURVIVAL, MANAGE EDEMA, AND TO REDUCE THE NEED FOR HOSPITALIZATION FOR HEART FAILURE
- U-2110 METHOD FOR CHRONIC WEIGHT MANAGEMENT IN PATIENTS WITH MODERATE RENAL IMPAIRMENT WHO ARE OBESE, OR OVERWEIGHT AND HAVE AT LEAST ONE WEIGHT RELATED COMORBID CONDITION
- U-2111 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIMS 1-5 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIMS 1-5
- U-2112 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIM 6 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 6
- U-2113 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIM 7 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 7
- U-2114 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIM 9 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 9
- U-2115 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIM 10 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 10
- U-2116 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIM 12 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 12
- U-2117 ADJUNCT TO DIET AND EXERCISE TO TREAT GLUCOSE INTOLERANCE IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIMS 14-15 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIMS 14-15
- U-2118 ADJUNCT TO DIET AND EXERCISE TO TREAT GLUCOSE INTOLERANCE IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIMS 16-18 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIMS 16-18
- U-2119 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN

PATENT AND EXCLUSIVITY TERMS

ADB 143 of 225

PATENT USE

- SECRETAGOGUE AS RECITED IN CLAIM 19 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 19
- U-2120 TREATMENT OF PATIENTS 18 YEARS OF AGE AND OLDER WITH COMPLICATED URINARY TRACT INFECTIONS CAUSED BY SUSCEPTIBLE MICROORGANISMS
- U-2121 TREATMENT OF PARTIAL-ONSET SEIZURES IN A PATIENT SUFFERING FROM OR SUSCEPTIBLE TO ABSENCE SEIZURES
- U-2122 USE FOR REDUCING EXACERBATIONS OF CHRONIC OBSTRUCTIVE PULMONARY DISEASE
- U-2123 TREATMENT OF PARTIAL-ONSET SEIZURES IN PATIENTS WITH EPILEPSY WHO HAVE BEEN PREVIOUSLY TREATED WITH OXCARBAZEPINE
- U-2124 TREATMENT OF ADULT PATIENTS WITH RELAPSED FOLLICULAR LYMPHOMA WHO HAVE RECEIVED AT LEAST TWO PRIOR SYSTEMIC THERAPIES
- U-2125 THE TREATMENT OF AN INFLAMMATORY DISORDER OF THE RESPIRATORY TRACT BY ONCE-PER-DAY ADMINISTRATION OF A PHARMACEUTICAL FORMULATION COMPRISING FLUTICASONE FUROATE AND A LONG-ACTING BETA2 ADRENORECEPTOR AGONIST
- U-2126 USE OF FLUTICASONE FUROATE FOR THE TREATMENT OF AN INFLAMMATORY OR ALLERGIC CONDITIONS, INCLUDING CHRONIC OBSTRUCTIVE PULMONARY DISEASE
- U-2127 INDICATED FOR THE LONG-TERM, ONCE-DAILY, MAINTENANCE TREATMENT OF PATIENTS WITH CHRONIC OBSTRUCTIVE PULMONARY DISEASE (COPD), INCLUDING CHRONIC BRONCHITIS AND/OR EMPHYSEMA
- U-2128 METHOD OF INHIBITING THE BINDING OF ACETYLCHOLINE TO AN ACETYLCHOLINE RECEPTOR IN THE RESPIRATORY TRACT OF A HUMAN, COMPRISING CONTACTING THE RECEPTOR WITH AN EFFECTIVE AMOUNT OF UMECLIDINIUM, VIA INHALATION
- U-2129 METHOD OF INHIBITING THE BINDING OF ACETYLCHOLINE TO AN ACETYLCHOLINE RECEPTOR IN THE RESPIRATORY TRACT OF A HUMAN, COMPRISING CONTACTING THE RECEPTOR WITH AN EFFECTIVE AMOUNT OF UMECLIDINIUM, VIA TOPICAL APPLICATION
- U-2130 TREATMENT OF PARTIAL ONSET SEIZURES IN PATIENTS WITH EPILEPSY AGED 16 YEARS AND OLDER WITH EPILEPSY
- U-2131 REDUCTION OF ELEVATED INTRAOCULAR PRESSURE (IOP) IN PATIENTS WITH GLAUCOMA OR OCULAR HYPERTENSION, WITH COMPARABLE EFFICACY, AND A REDUCTION IN SPECIFIED ADVERSE EVENTS, COMPARED TO BRIMONIDINE 0.2% TID
- U-2132 IN COMBINATION WITH FULVESTRANT FOR THE TREATMENT OF WOMEN WITH HORMONE RECEPTOR (HR)-POSITIVE, HUMAN EPIDERMAL GROWTH FACTOR RECEPTOR 2 (HER2)-NEGATIVE ADVANCED OR METASTATIC BREAST CANCER WITH DISEASE PROGRESSION FOLLOWING ENDOCRINE THERAPY
- U-2133 METHOD OF DELIVERING FLUTICASONE PROPIONATE TO A NASAL AIRWAY
- U-2134 THE TREATMENT OF CHRONIC OBSTRUCTIVE PULMONARY DISEASE BY ONCE-PER-DAY ADMINISTRATION OF A PHARMACEUTICAL FORMULATION COMPRISING FLUTICASONE FUROATE AND A LONG-ACTING BETA2 ADRENORECEPTOR
- U-2135 AS MONOTHERAPY FOR THE TREATMENT OF ADULT PATIENTS WITH HR-POSITIVE, HER2-NEGATIVE ADVANCED OR METASTATIC BREAST CANCER WITH DISEASE PROGRESSION FOLLOWING ENDOCRINE THERAPY AND PRIOR CHEMOTHERAPY IN THE METASTATIC SETTING
- U-2136 TREATMENT OF NEUROPATHIC PAIN ASSOCIATED WITH DIABETIC PERIPHERAL NEUROPATHY
- U-2137 TREATMENT OF POSTHERPETIC NEURALGIA
- U-2138 TOPICAL TREATMENT OF ACTINIC KERATOSIS OF THE FACE OR SCALP USING MORE THAN ONE TREATMENT COURSE OF INGENOL MEBUTATE
- U-2139 TREATMENT OF TYPE 2 DIABETES MELLITUS IN COMBINATION WITH EXENATIDE
- U-2140 METHOD OF TREATING PARTIAL ONSET SEIZURES IN PATIENTS 4 YEARS OF AGE AND OLDER
- U-2141 TREATMENT OF CHRONIC HEPATITIS C VIRUS (HCV) GENOTYPE 1, 2, 3, 4, 5, OR 6
- U-2142 REDUCTION IN THE RISK OF RECURRENCE OF DEEP VEIN THROMBOSIS (DVT) AND/OR PULMONARY EMBOLISM (PE) IN PATIENTS AT CONTINUED RISK FOR RECURRENT DVT AND/OR AFTER COMPLETION OF INITIAL TREATMENT LASTING AT LEAST 6 MONTHS
- U-2143 AFTER COMPLETION OF INITIAL TREATMENT LASTING AT LEAST 6 MONTHS, TO REDUCE THE RISK OF RECURRENCE OF DEEP VEIN THROMBOSIS AND/OR PULMONARY EMBOLISM IN CERTAIN PATIENTS WITH ONCE DAILY, RAPID-RELEASE TABLET ADMINISTERED FOR AT LEAST 5 CONSECUTIVE DAYS
- U-2144 REDUCTION OF INTRAOCULAR PRESSURE (IOP) IN PATIENTS WITH OPEN-ANGLE GLAUCOMA OR OCULAR HYPERTENSION
- U-2145 TREATMENT OF ADULT PATIENTS WITH MANTLE CELL LYMPHOMA WHO HAVE RECEIVED AT LEAST ONE PRIOR THERAPY

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-2146 IMPROVEMENT IN GLYCEMIC CONTROL IN DIABETES MELLITUS PATIENTS BY USE OF A PEN INJECTOR WITH A ROTATING DRIVE SLEEVE

U-2147 TREATMENT OF NON-24 HOUR SLEEP-WAKE DISORDER BY ORALLY ADMINISTERING 20MG OF TASIMELTEON ONCE DAILY BEFORE BEDTIME

U-2148 A METHOD OF PROVIDING NITRIC OXIDE THERAPY TO A PATIENT BY MEASURING AND DISPLAYING AN INDICATION OF THE CALCULATED DELIVERY CONCENTRATION OF NITRIC OXIDE AS COMPARED TO THE DESIRED DELIVERY CONCENTRATION OF NITRIC OXIDE

U-2149 TREATMENT OF NON-24 HOUR SLEEP-WAKE DISORDER BY ADMINISTERING TASIMELTEON

U-2150 TREATMENT OF CHRONIC GRAFT-VERSUS-HOST DISEASE

U-2151 METHOD OF TREATING PAIN OR INFLAMMATION WITH AN INJECTABLE CONTROLLED OR SUSTAINED RELEASE FORMULATION OF TRIAMCINOLONE ACETONIDE

U-2152 TREATMENT OF PAIN ASSOCIATED WITH IRRITABLE BOWEL SYNDROME WITH DIARRHEA (IBS-D) WITH VIBERZI (ELUXADOLINE)

U-2153 REDUCING FASTING PLASMA GLUCOSE IN A HUMAN IN NEED THEREOF USING A SUSTAINED-RELEASE COMPOSITION CONTAINING EXENDIN-4

U-2154 REDUCING FASTING PLASMA GLUCOSE IN A HUMAN IN NEED THEREOF USING A SUSTAINED-RELEASE COMPOSITION CONTAINING EXENDIN-4

U-2155 REDUCING BODY WEIGHT IN A HUMAN IN NEED THEREOF USING A SUSTAINED-RELEASE COMPOSITION CONTAINING EXENDIN-4

U-2156 REDUCING HBA1C IN A HUMAN IN NEED THEREOF USING A SUSTAINED-RELEASE COMPOSITION CONTAINING EXENDIN-4

U-2157 TREATING TYPE 2 DIABETES MELLITUS BY STIMULATING INSULIN RELEASE

U-2158 DECREASING GASTRIC MOTILITY OR DELAYING GASTRIC EMPTYING BY USING A SUSTAINED-RELEASE COMPOSITION

U-2159 TREATMENT OF CHRONIC LYMPHOCYTIC LEUKEMIA/SMALL LYMPHOCYTIC LYMPHOMA

U-2160 MANAGEMENT OF OSTEOARTHRITIS PAIN BY ADMINISTERING 5 MG OF MELOXICAM

U-2161 TREATMENT OF NAUSEA AND VOMITING, INCLUDING THE PREVENTION OF ACUTE AND DELAYED NAUSEA AND VOMITING ASSOCIATED WITH INITIAL AND REPEAT COURSES OF HIGHLY OR MODERATELY EMETOGENIC CANCER CHEMOTHERAPY

U-2162 FOR CLEANSING THE LARGE INTESTINE AS A PREPARATION FOR COLONOSCOPY

U-2163 TREATMENT OF HR-POSITIVE, HUMAN EPIDERMAL GROWTH FACTOR RECEPTOR 2 (HER-2)-NEGATIVE ADVANCED OR METASTATIC BREAST CANCER IN COMBINATION WITH PALBOCICLIB OR ABEMACICLIB IN WOMEN WITH DISEASE PROGRESSION AFTER ENDOCRINE THERAPY

U-2164 ZELBORAF IS INDICATED FOR THE TREATMENT OF PATIENTS WITH ERDHEIM-CHESTER DISEASE WITH BRAF V600 MUTATION

U-2165 MANAGEMENT OF OSTEOARTHRITIS PAIN BY ADMINISTERING 10 MG OF MELOXICAM

U-2166 TREATMENT OF MAJOR DEPRESSIVE EPISODES ASSOCIATED WITH BIPOLAR I DISORDER

U-2167 METHOD OF USING A TABLET EMBEDDED WITH A SENSOR THAT COMMUNICATES INFORMATION VIA A SIGNAL THROUGH THE BODY OF A PATIENT TO A RECEIVER

U-2168 METHOD OF USING A LOGIC CIRCUIT TO STABILIZE BATTERY VOLTAGE SUPPLIED TO A SENSOR EMBEDDED WITH A TABLET AND THAT COMMUNICATES INFORMATION VIA A SIGNAL THROUGH THE BODY OF A PATIENT TO A RECEIVER

U-2169 METHOD OF USING A RECEIVER TO IDENTIFY A SIGNAL FROM A TABLET EMBEDDED WITH A SENSOR THAT COMMUNICATES INFORMATION THROUGH THE BODY OF A PATIENT

U-2170 METHOD OF USING A RECEIVER TO RECEIVE A SIGNAL FROM A TABLET EMBEDDED WITH A SENSOR THAT COMMUNICATES INFORMATION THROUGH THE BODY OF A PATIENT

U-2171 ADJUVANT TREATMENT OF ADULT PATIENTS AT HIGH RISK OF RECURRENT RCC FOLLOWING NEPHRECTOMY

U-2172 METHOD TO TREAT SEVERE ALLERGIC EMERGENCIES IN PATIENTS WEIGHING 7.5 TO 15 KG (16.5 TO 33 LBS)

U-2173 TREATING OPIOID DEPENDENCE BY ADMINISTERING BUPRENORPHINE

U-2174 TREATING OPIOID DEPENDENCY BY ADMINISTERING BUPRENORPHINE ONCE PER MONTH

U-2175 TREATING OPIOID DEPENDENCY BY ADMINISTERING BUPRENORPHINE ONCE MONTHLY

U-2176 TREATING OPIOID ADDICTION BY ADMINISTERING BUPRENORPHINE

U-2177 TREATING OPIOID ADDICTION BY SUBCUTANEOUS INJECTION OF BUPRENORPHINE

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-2178 TREATING OPIOID ADDICTION BY ADMINISTERING BUPRENORPHINE COMPOSITION WITH 28 DAY DOSE DURATION

U-2179 IN SITU FORMATION OF SOLID BUPRENORPHINE COMPOSITION

U-2180 TREATING ADDICTION WITH 100 MG OR 300 MG DOSE OF BUPRENORPHINE

U-2181 TREATING OPIOID DEPENDENCY BY SUBCUTANEOUSLY ADMINISTERING BUPRENORPHINE

U-2182 IMPROVEMENT OF GLYCEMIC CONTROL IN TYPE 2 DIABETES MELLITUS PATIENTS

U-2183 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 1 AND 13

U-2184 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 1, 13, AND 14

U-2185 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING A DOSAGE FORM COMPRISING MICRONIZED BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 15 AND 27

U-2186 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING A DOSAGE FORM COMPRISING MICRONIZED BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 15, 27, AND 28

U-2187 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 29 AND 39

U-2188 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 29, 39, AND 40

U-2189 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 41 AND 52

U-2190 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 41, 52, AND 53

U-2191 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 54 AND 64

U-2192 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 54, 64, AND 65

U-2193 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING A DOSAGE FORM COMPRISING MICRONIZED BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 66 AND 75

U-2194 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING A DOSAGE FORM COMPRISING MICRONIZED BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 66, 75, AND 76

U-2195 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 77 AND 87

U-2196 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 77, 87, AND 88

U-2197 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 89 AND 99

U-2198 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 89, 99, AND 100

U-2199 TREATMENT OF SCHIZOPHRENIA WITH IMPROVEMENT IN ATTENTION FUNCTION IN SCHIZOPHRENIA

U-2200 COMBINATION TREATMENT WITH INSULIN GLARGINE WITH OR WITHOUT METFORMIN FOR IMPROVEMENT OF GLYCEMIC CONTROL IN TYPE 2 DIABETES MELLITUS PATIENTS

U-2201 TREATMENT OF BIPOLAR DEPRESSION WITH IMPROVEMENT IN ATTENTION FUNCTION IN

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

BIPOLAR DISORDER

- U-2202 OZEMPIC IS INDICATED AS AN ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS
- U-2203 A METHOD OF PROVIDING A SUBJECT WITH A THERAPEUTICALLY EFFECTIVE AMOUNT OF RACEMIC METHYLPHENIDATE BY ORALLY ADMINISTERING TO SAID SUBJECT A SINGLE METHYLPHENIDATE EXTENDED RELEASE CHEWABLE TABLET AS CLAIMED
- U-2204 TREATING PATIENTS WITH ACUTE PROMYELOCYTIC LEUKEMIA (APL) WHO ARE REFRACTORY TO, OR HAVE RELAPSED FROM, RETINOID AND ANTHRACYCLINE CHEMOTHERAPY, AND WHOSE APL IS CHARACTERIZED BY THE PRESENCE OF THE T(15;17) TRANSLOCATION OR PML/RAR-ALPHA GENE EXPRESSION
- U-2205 TREATMENT OF SEBORRHEIC KERATOSES THAT ARE RAISED
- U-2206 TREATING OPIOID DEPENDENCY BY ADMINISTERING BUPRENORPHINE
- U-2207 TREATING ADDICTION BY SUBCUTANEOUS INJECTION OF BUPRENORPHINE
- U-2208 TREATING ADDICTION BY ONCE PER MONTH ADMINISTRATION OF BUPRENORPHINE
- U-2209 TREATING OPIOID ADDICTION BY ADMINISTERING BUPRENORPHINE ONCE PER MONTH
- U-2210 TREATING OPIOID ADDICTION BY 100 MG OR 300 MG DOSE BUPRENORPHINE
- U-2211 TREATING OPIOID ADDICTION BY ADMINISTRATION OF BUPRENORPHINE
- U-2212 REDUCING FASTING PLASMA GLUCOSE IN A HUMAN IN NEED THEREOF IN COMBINATION WITH A SUSTAINED-RELEASE COMPOSITION CONTAINING EXENDIN-4
- U-2213 REDUCING HBA1C IN A HUMAN IN NEED THEREOF IN COMBINATION WITH A SUSTAINED-RELEASE COMPOSITION CONTAINING EXENDIN-4
- U-2214 AN ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES
- U-2215 ERTUGLIFLOZIN IN COMBINATION WITH SITAGLIPTIN AND IN FURTHER COMBINATION WITH METFORMIN AS AN ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS
- U-2216 ERTUGLIFLOZIN AND SITAGLIPTIN IN COMBINATION AS AN ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS
- U-2217 TREATING HIGH OUTPUT SHOCK WITH ANGIOTENSIN II BY INCREASING MEAN ARTERIAL PRESSURE IN PATIENTS TREATED WITH CATECHOLAMINES AND REDUCING CATECHOLAMINE USE
- U-2218 MAINTAINING MEAN ARTERIAL PRESSURE OF ABOUT 65 MMHG OR HIGHER WITH ANGIOTENSIN II IN SHOCK PATIENTS TREATED WITH CATECHOLAMINES AND REDUCING CATECHOLAMINE USE
- U-2219 TREATMENT OF CHRONIC SMALL LYMPHOCYTIC LEUKEMIA
- U-2220 A METHOD FOR THE DIAGNOSIS OF ADULT GROWTH HORMONE DEFICIENCY BY MEASURING THE LEVEL OF GROWTH HORMONE AFTER ORAL ADMINISTRATION OF MACIMORELIN
- U-2221 TREATING REFRACTORY HYPOTENSION WITH ABOUT 20 NG/KG/MIN ANGIOTENSIN II IN A PATIENT RECEIVING VASOPRESSOR
- U-2222 RELIEVES REDNESS OF THE EYE DUE TO MINOR EYE IRRITATIONS
- U-2223 METHOD OF TREATING ANGINA PECTORIS
- U-2224 TREATMENT OF DYSKINESIA AND INCREASING ON TIME WITHOUT TROUBLESOME DYSKINESIA IN PATIENTS WITH PARKINSON'S DISEASE RECEIVING LEVODOPA-BASED THERAPY, WITH OR WITHOUT CONCOMITANT DOPAMINERGIC MEDICATIONS
- U-2225 METHOD OF ADMINISTERING A LOCAL ANESTHETIC TO THE MUCOUS MEMBRANES IN PATIENTS WITH HEPATIC IMPAIRMENT
- U-2226 METHOD OF ADMINISTERING A LOCAL ANESTHETIC TO THE MUCOUS MEMBRANES IN PATIENTS WITH RENAL IMPAIRMENT
- U-2227 METHOD OF ADMINISTERING A LOCAL ANESTHETIC TO THE MUCOUS MEMBRANES IN GERIATRIC PATIENTS
- U-2228 TREATMENT OF SMALL LYMPHOCYTIC LEUKEMIA
- U-2229 IN COMBINATION WITH TRETINOIN, TREATING ADULTS AND PEDIATRIC PATIENTS 1 YEAR AND OLDER WITH NEWLY-DIAGNOSED LOW-RISK ACUTE PROMYELOCYTIC LEUKEMIA (APL) CHARACTERIZED BY THE PRESENCE OF THE T(15;17) TRANSLOCATION OR PML/RAR-A GENE EXPRESSION
- U-2230 IRRITABLE BOWEL SYNDROME WITH CONSTIPATION
- U-2231 TREATING REFRACTORY HYPOTENSION WITH ABOUT 5 NG/KG/MIN TO ABOUT 20 NG/KG/MIN

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

ANGIOTENSIN II IN A PATIENT RECEIVING VASOPRESSOR

- U-2232 TREATMENT OF PSORIATIC ARTHRITIS USING A DOSAGE TITRATION SCHEDULE
- U-2233 TREATMENT OF PSORIATIC ARTHRITIS WITH APREMILAST USING A DOSAGE TITRATION SCHEDULE AND A SECOND ACTIVE AGENT
- U-2234 USE OF IVACAFTOR FOR TREATING CYSTIC FIBROSIS IN A PATIENT WITH A MILD TO MODERATE CF PHENOTYPE WITH AT LEAST ONE MUTATION IN THE CFTR GENE THAT IS RESPONSIVE TO IVACAFTOR BASED ON CLINICAL AND/OR IN VITRO ASSAY DATA
- U-2235 USE IN COMBINATION WITH PREDNISONE FOR THE TREATMENT OF PATIENTS WITH METASTATIC HIGH-RISK CASTRATION-SENSITIVE PROSTATE CANCER
- U-2236 REDUCING THE RISK OF PRETERM BIRTH IN WOMEN WITH A SINGLETON PREGNANCY WHO HAVE A HISTORY OF SINGLETON SPONTANEOUS PRETERM BIRTH
- U-2237 TREATMENT OF NON-METASTATIC, CASTRATION-RESISTANT PROSTATE CANCER (NM-CRPC)
- U-2238 METHOD OF IMPROVING GLYCEMIC CONTROL IN PATIENTS WITH DIABETES MELLITUS BY ADMINISTERING A MIXTURE OF INSULIN DEGLUDEC AND INSULIN ASPART DURING OR AROUND THE TIME OF THE LARGEST MEAL OF THE DAY
- U-2239 REDUCTION OF ELEVATED INTRAOCULAR PRESSURE (IOP) IN PATIENTS WITH GLAUCOMA OR OCULAR HYPERTENSION, WITH A REDUCTION IN SPECIFIED ADVERSE EVENTS, COMPARED TO BRIMONIDINE 0.2% TID
- U-2240 REDUCTION OF ELEVATED INTRAOCULAR PRESSURE (IOP) IN PATIENTS WITH GLAUCOMA OR OCULAR HYPERTENSION, WITH COMPARABLE EFFICACY TO BRIMONIDINE 0.2% TID
- U-2241 TREATMENT OF SMALL LYMPHOCYTIC LYMPHOMA WITH 17P DELETION
- U-2242 TREATMENT OF CHRONIC LYMPHOCYTIC LEUKEMIA WITH 17P DELETION
- U-2243 TREATMENT OF CHRONIC LYMPHOCYTIC LEUKEMIA/SMALL LYMPHOCYTIC LYMPHOMA WITH 17P DELETION
- U-2244 A METHOD OF TREATING BACTERIAL INFECTIONS IN HOSPITAL-ACQUIRED BACTERIAL PNEUMONIA AND VENTILATOR-ASSOCIATED BACTERIAL PNEUMONIA (HABP/VABP) PATIENTS COMPRISING ADMINISTERING A BACTERICIDALLY EFFECTIVE AMOUNT OF AVIBACTAM SODIUM
- U-2245 A METHOD OF TREATING A BACTERIAL INFECTION IN HOSPITAL-ACQUIRED BACTERIAL PNEUMONIA AND VENTILATOR-ASSOCIATED BACTERIAL PNEUMONIA (HABP/VABP) PATIENTS COMPRISING ADMINISTERING AN EFFECTIVE AMOUNT OF AVIBACTAM SODIUM
- U-2246 TEZACAFTOR AND IVACAFTOR FOR THE TREATMENT OF CYSTIC FIBROSIS IN PATIENTS WHO ARE HOMOZYGOUS FOR THE F508DEL MUTATION OR HAVING AT LEAST ONE CFTR GENE MUTATION THAT IS RESPONSIVE TO TEZACAFTOR/IVACAFTOR BASED ON IN VITRO DATA AND/OR CLINICAL EVIDENCE
- U-2247 TEZACAFTOR AND IVACAFTOR FOR THE TREATMENT OF PATIENTS WITH A MILD TO MODERATE CLINICAL PHENOTYPE OF CYSTIC FIBROSIS HAVING AT LEAST ONE CFTR GENE MUTATION THAT IS RESPONSIVE TO TEZACAFTOR/IVACAFTOR BASED ON IN VITRO DATA AND/OR CLINICAL EVIDENCE
- U-2248 TEZACAFTOR AND IVACAFTOR FOR THE TREATMENT OF CYSTIC FIBROSIS IN PATIENTS WHO ARE HOMOZYGOUS FOR THE F508DEL MUTATION OR HETEROZYGOUS FOR THE F508DEL MUTATION AND A SECOND MUTATION THAT IS RESPONSIVE TO TEZACAFTOR/IVACAFTOR
- U-2249 MANAGEMENT OF ACUTE PAIN SEVERE ENOUGH TO REQUIRE AN OPIOID ANALGESIC AND FOR WHICH ALTERNATIVE TREATMENTS ARE INADEQUATE
- U-2250 DETECTION OF CARCINOMA IN THE BLADDER BY PHOTODYNAMIC CYSTOSCOPY
- U-2251 IN COMBINATION WITH AN AROMATASE INHIBITOR AS INITIAL ENDOCRINE-BASED THERAPY FOR THE TREATMENT OF POSTMENOPAUSAL WOMEN WITH HORMONE RECEPTOR (HR)-POSITIVE, HUMAN EPIDERMAL GROWTH FACTOR RECEPTOR 2 (HER2)-NEGATIVE ADVANCED OR METASTATIC BREAST CANCER
- U-2252 THE TREATMENT OF ACUTE OTITIS EXTERNA IN PATIENTS 6 MONTHS OF AGE AND OLDER DUE TO PSEUDOMONAS AERUGINOSA AND STAPHYLOCOCCUS AUREUS
- U-2253 PROPHYLACTIC TREATMENT OF NAUSEA AND VOMITING, INCLUDING PREVENTION OF ACUTE AND DELAYED NAUSEA AND VOMITING ASSOCIATED CHEMOTHERAPY
- U-2254 USE OF POMALIDOMIDE WITH DEXAMETHASONE FOR PATIENTS WITH MULTIPLE MYELOMA AFTER AT LEAST TWO PRIOR THERAPIES INCLUDING LENALIDOMIDE AND A PROTEASOME INHIBITOR AND DEMONSTRATED DISEASE PROGRESSION ON OR WITHIN 60 DAYS OF COMPLETING THE LAST THERAPY
- U-2255 TREATING SECONDARY HYPERPARATHYROIDISM IN CHRONIC KIDNEY DISEASE WITH SUSTAINED RELEASE 25-HYDROXYVITAMIN D TO REDUCE THE PATIENT'S SERUM PARATHYROID HORMONE

PATENT AND EXCLUSIVITY TERMS

ADB 148 of 225

PATENT USE

LEVEL AND THE SUSTAINED RELEASE IS OVER AT LEAST 10 HOURS

- U-2256 TREATING SECONDARY HYPERPARATHYROIDISM IN CHRONIC KIDNEY DISEASE WITH SUSTAINED RELEASE 25-HYDROXYVITAMIN D TO REDUCE THE PATIENT'S SERUM PARATHYROID HORMONE LEVEL AND CMAX IS REDUCED COMPARED TO BOLUS IV INJECTION AND IMMEDIATE-RELEASE, ORAL DOSING
- U-2257 TREATING SHPT IN CKD WITH SUSTAINED RELEASE CALCIFEDIOL TO REDUCE SERUM PARATHYROID HORMONE LEVEL AND CHANGE IN SERUM CONCENTRATION OF CALCIFEDIOL IN DOSE INTERVAL IS REDUCED COMPARED TO BOLUS IV INJECTION AND IMMEDIATE-RELEASE, ORAL DOSING
- U-2258 TREATING SECONDARY HYPERPARATHYROIDISM IN CKD WITH SUSTAINED RELEASE CALCIFEDIOL TO REDUCE THE PATIENT'S SERUM PARATHYROID HORMONE LEVEL AND CMAX24HR/C24HR IS REDUCED COMPARED TO BOLUS IV INJECTION AND IMMEDIATE-RELEASE, ORAL DOSING
- U-2259 TREATING SECONDARY HYPERPARATHYROIDISM IN CKD WITH SUSTAINED RELEASE CALCIFEDIOL TO REDUCE THE PATIENT'S SERUM PARATHYROID HORMONE LEVEL AND TMAX IS INCREASED COMPARED TO BOLUS IV INJECTION AND IMMEDIATE-RELEASE, ORAL DOSING
- U-2260 METHOD OF REDUCING THE RISK OF PERIPROCEDURAL MYOCARDIAL INFARCTION, AND STENT THROMBOSIS IN A PATIENT UNDERGOING PCI BY ADMINISTERING INTRAVENOUSLY 30 UG/KG BOLUS BEFORE PCI AND THEN A CONTINUOUS INFUSION
- U-2261 MODIFIED DOSING REGIMEN FOR THE MANAGEMENT OF MILD TO MODERATE PAIN OR MANAGEMENT OF MODERATE TO SEVERE PAIN AS AN ADJUNCT TO OPIOID ANALGESICS
- U-2262 MODIFIED DOSING REGIMEN FOR THE REDUCTION OF FEVER
- U-2263 MODIFIED DOSING REGIMEN FOR THE MANAGEMENT OF MODERATE TO SEVERE PAIN WITH ADJUNCTIVE OPIOID ANALGESICS
- U-2264 METHODS OF TREATING PAIN, INFLAMMATION, FEVER, PATENT DUCTUS ARTERIOSIS WITH AQUEOUS COMPOSITION
- U-2265 PREVENTION OF NAUSEA AND VOMITING ASSOCIATED WITH HEC AND MEC IN ADULT AND PEDIATRIC PATIENTS
- U-2266 METHODS OF MAKING AQUEOUS COMPOSITION AND TREATING PAIN, INFLAMMATION, FEVER, PATENT DUCTUS ARTERIOSIS WITH AQUEOUS COMPOSITION
- U-2267 METHOD FOR RELIEVING THE PAIN ASSOCIATED WITH POST-HERPETIC NEURALGIA
- U-2268 DISCONTINUING A STRONG CYP1A2 INDUCER TO AVOID REDUCED PIRFENIDONE EFFICACY AND THEN ADMINISTERING PIRFENIDONE
- U-2269 DOSAGE MODIFICATION FOLLOWING GRADE 2 ABNORMALITY IN LIVER FUNCTION BIOMARKER AFTER PIRFENIDONE ADMINISTRATION, BY ADMINISTERING SUB-2400MG/DAY DOSE THEN FULL DAILY DOSE IN TREATMENT OF IPF
- U-2270 DOSAGE MODIFICATION FOLLOWING GRADE 2 ABNORMALITY IN LIVER FUNCTION BIOMARKER AFTER PIRFENIDONE ADMINISTRATION, BY DISCONTINUING PIRFENIDONE UNTIL BIOMARKERS ARE WITHIN NORMAL LIMITS, THEN SUB-2400MG/DAY DOSE, THEN FULL DAILY DOSE IN TREATMENT OF IPF
- U-2271 THERAPEUTIC TREATMENT OF PATIENTS WITH CASTRATION-RESISTANT PROSTATE CANCER, SYMPTOMATIC BONE METASTASES AND NO KNOWN VISCERAL METASTATIC DISEASE
- U-2272 TREATMENT OF NASAL POLYPS IN PATIENTS ≥ 18 YEARS OF AGE WHO HAVE HAD ETHMOID SINUS SURGERY USING A CORTICOSTEROID-ELUTING (MOMETASONE FUROATE) IMPLANT
- U-2273 A METHOD FOR TREATING EPITHELIAL OVARIAN, FALLOPIAN TUBE, OR PRIMARY PERITONEAL CANCER, WHEREIN THE CANCER IS ASSOCIATED WITH A DELETERIOUS BRCA MUTATION
- U-2274 MAINTAINING SERUM 25-HYDROXYVITAMIN D AT A LEVEL OF AT LEAST 30 NG/ML WITH ORAL, SUSTAINED RELEASE 25-HYDROXYVITAMIN D
- U-2275 TREATING CYSTIC FIBROSIS PATIENTS AGES 12 AND OLDER, WHO ARE HOMOZYGOUS FOR F508DEL OR HAVE AT LEAST 1 CFTR GENE MUTATION RESPONSIVE TO TEZACAFTOR/IVACAFTOR, WITH TEZACAFTOR AND A SOLID COMPOSITION COMPRISING AMORPHOUS (<30% CRYSTALLINE) IVACAFTOR
- U-2276 METHOD OF TREATING CYSTIC FIBROSIS IN A PATIENT AGE 6 OR OLDER HOMOZYGOUS FOR THE F508DEL MUTATION IN THE CFTR GENE USING LUMACAFTOR AND A SOLID COMPOSITION COMPRISING AMORPHOUS (LESS THAN ABOUT 30% CRYSTALLINE) IVACAFTOR
- U-2277 IMPROVEMENT IN GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS INADEQUATELY CONTROLLED BY LIXISENATIDE
- U-2278 IMPROVEMENT IN GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS INADEQUATELY CONTROLLED BY LIXISENATIDE IN COMBINATION WITH METFORMIN

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

- U-2279 IMPROVEMENT IN GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS INADEQUATELY CONTROLLED BY LIXISENATIDE IN COMBINATION WITH METFORMIN AND A SECOND ORAL ANTIDIABETIC DRUG
- U-2280 ADJUNCTIVE TREATMENT OF PATIENTS WITH TSC-ASSOCIATED PARTIAL-ONSET SEIZURES
- U-2281 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIM 1 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 1
- U-2282 ADJUNCT TO DIET AND EXERCISE TO TREAT GLUCOSE INTOLERANCE IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIM 2 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 2
- U-2283 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIMS 3-7 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIMS 3-7
- U-2284 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIM 8 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 8
- U-2285 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIM 11 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 11
- U-2286 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIM 14 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 14
- U-2287 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIMS 16-19 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIMS 16-19
- U-2288 TREATMENT OF TYPE 2 DIABETES MELLITUS WITH EXENATIDE AS AN ADD-ON TO BASIL INSULIN OR BASAL INSULIN PLUS METFORMIN THERAPY
- U-2289 TREATMENT OF PATIENTS WITH METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) WHOSE TUMORS HAVE EPIDERMAL GROWTH FACTOR RECEPTOR (EGFR) EXON 19 DELETIONS OR EXON 21L858R MUTATIONS
- U-2290 METHOD OF TREATING TYPE 2 DIABETES MELLITUS IN A PATIENT WITH RENAL IMPAIRMENT (45 ML/MIN/1.73 M² ≤ EGFR < 60 ML/MIN/1.73 M²) BY ONCE DAILY ADMINISTRATION OF 10 MG OR 25 MG OF EMPAGLIFLOZIN
- U-2291 REDUCTION OF THROMBOTIC CARDIOVASCULAR EVENTS IN PATIENTS WITH A HISTORY OF MYOCARDIAL INFARCTION (MI) OR WITH PERIPHERAL ARTERIAL DISEASE (PAD)
- U-2292 METHOD OF REDUCING THE RISK OF CARDIOVASCULAR DEATH IN ADULT PATIENTS WITH TYPE 2 DIABETES MELLITUS AND CARDIOVASCULAR DISEASE BY ONCE DAILY ADMINISTRATION OF 10 MG OR 25 MG OF EMPAGLIFLOZIN
- U-2293 USE IN COMBINATION WITH DEXAMETHASONE IN ADULTS FOR THE PREVENTION OF ACUTE AND DELAYED NAUSEA AND VOMITING ASSOCIATED WITH INITIAL AND REPEAT COURSES OF CANCER CHEMOTHERAPY, INCLUDING, BUT NOT LIMITED TO, HIGHLY EMETOGENIC CHEMOTHERAPY
- U-2294 TREATMENT OF THROMBOCYTOPENIA IN ADULT PATIENTS WITH CHRONIC IMMUNE THROMBOCYTOPENIA (ITP) WHO HAVE HAD AN INSUFFICIENT RESPONSE TO A PREVIOUS TREATMENT
- U-2295 TREATMENT OF PARTIAL-ONSET SEIZURES IN PATIENTS 4 YEARS OF AGE AND OLDER
- U-2296 TAFINLAR IS INDICATED, IN COMBINATION WITH TRAMETINIB, FOR THE ADJUVANT TREATMENT OF PATIENTS WITH MELANOMA WITH BRAF V600E OR V600K MUTATIONS, AS DETECTED BY AN FDA-APPROVED TEST, AND INVOLVEMENT OF LYMPH NODE(S), FOLLOWING COMPLETE RESECTION
- U-2297 IMPROVEMENT OF GLYCEMIC CONTROL IN TYPE 2 DIABETES PATIENTS BY ADMINISTERING A STARTING DOSE OF 10 MCG FOR 14 DAYS AND INCREASING TO A MAINTENANCE DOSE OF 20 MCG ON DAY 15
- U-2298 TAFINLAR IS INDICATED, IN COMBINATION WITH TRAMETINIB, FOR THE TREATMENT OF PATIENTS WITH LOCALLY ADVANCED OR METASTATIC ANAPLASTIC THYROID CANCER (ATC)

PATENT AND EXCLUSIVITY TERMS

ADB 150 of 225

PATENT USE

WITH BRAF V600E MUTATION AND WITH NO SATISFACTORY LOCOREGIONAL TREATMENT OPTIONS

U-2299 TAFINLAR IS INDICATED AS A SINGLE AGENT FOR THE TREATMENT OF PATIENTS WITH UNRESECTABLE OR METASTATIC MELANOMA WITH BRAF V600E MUTATION

U-2300 USE IN COMBINATION WITH THE MUSCARINIC ANTAGONIST SOLIFENACIN SUCCINATE FOR THE TREATMENT OF OVERACTIVE BLADDER (OAB) WITH SYMPTOMS OF URGE URINARY INCONTINENCE, URGENCY, AND URINARY FREQUENCY

U-2301 USE IN COMBINATION WITH DEXAMETHASONE IN ADULTS FOR THE PREVENTION OF ACUTE AND DELAYED NAUSEA AND VOMITING ASSOCIATED WITH INITIAL AND REPEAT COURSES OF HIGHLY EMETOGENIC CANCER CHEMOTHERAPY

U-2302 MEKINIST IS INDICATED, IN COMBINATION WITH DABRAFENIB, FOR THE ADJUVANT TREATMENT OF PATIENTS WITH MELANOMA WITH BRAF V600E OR V600K MUTATIONS, AS DETECTED BY AN FDA-APPROVED TEST, AND INVOLVEMENT OF LYMPH NODE(S), FOLLOWING COMPLETE RESECTION

U-2303 MEKINIST IS INDICATED, IN COMBINATION WITH DABRAFENIB, FOR THE ADJUVANT TREATMENT OF PATIENTS WITH MELANOMA WITH BRAF V600E OR V600K MUTATIONS AND INVOLVEMENT OF LYMPH NODE(S), FOLLOWING COMPLETE RESECTION

U-2304 MEKINIST IS INDICATED, IN COMBINATION WITH DABRAFENIB, FOR THE TREATMENT OF PATIENTS WITH METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) WITH BRAF V600E MUTATION

U-2305 MEKINIST IS INDICATED, IN COMBINATION WITH DABRAFENIB, FOR THE TREATMENT OF PATIENTS WITH LOCALLY ADVANCED OR METASTATIC ANAPLASTIC THYROID CANCER (ATC) WITH BRAF V600E MUTATION AND WITH NO SATISFACTORY LOCOREGIONAL TREATMENT OPTIONS

U-2306 ONCE DAILY TOPICAL TREATMENT OF PERSISTENT FACIAL ERYTHEMA ASSOCIATED WITH ROSACEA IN ADULTS WITH 1% OXYMETAZOLINE HYDROCHLORIDE CREAM

U-2307 TREATMENT OF AUTOSOMAL DOMINANT POLYCYSTIC KIDNEY DISEASE

U-2308 TREATMENT OF ADULT PATIENTS WITH SHORT BOWEL SYNDROME WHO ARE DEPENDENT ON PARENTERAL SUPPORT

U-2309 USE IN THE TREATMENT OF MAJOR DEPRESSIVE DISORDER TO IMPROVE PROCESSING SPEED, AN ASPECT OF COGNITIVE FUNCTION

U-2310 FOR CLEANSING OF THE COLON IN PREPARATION FOR COLONOSCOPY IN ADULTS

U-2311 TREATMENT OF HYPERURICEMIA ASSOCIATED WITH GOUT IN PATIENTS WHO HAVE NOT ACHIEVED TARGET SERUM URIC ACID LEVELS WITH A XANTHINE OXIDASE INHIBITOR ALONE

U-2312 TREATMENT OF HYPERKALEMIA IN ADULTS

U-2313 METHOD OF REDUCING THE RISK OF CARDIOVASCULAR DEATH, NON-FATAL MYOCARDIAL INFARCTION, AND/OR NON-FATAL STROKE IN ADULTS WITH TYPE 2 DIABETES MELLITUS AND ESTABLISHED CARDIOVASCULAR DISEASE BY ADMINISTERING LIRAGLUTIDE

U-2314 TREATMENT OF THROMBOCYTOPENIA IN AN ADULT PATIENT WITH CHRONIC LIVER DISEASE WHO IS SCHEDULED TO UNDERGO A PROCEDURE USING DOPTelet

U-2315 TREATMENT OF MULTIPLE SCLEROSIS IN THE PEDIATRIC PATIENT POPULATION WITH 0.25 MG FINGOLIMOD

U-2316 TREATMENT OF DYSPAREUNIA

U-2317 TREATMENT OF A SYMPTOM OF VULVAR AND VAGINAL ATROPHY

U-2318 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 12 AND OLDER, WHO ARE HOMOZYGOUS FOR THE F508DEL MUTATION OR HAVE AT LEAST ONE CFTR GENE MUTATION THAT IS RESPONSIVE TO TEZACAFTOR/IVACAFTOR, WITH TEZACAFTOR AND IVACAFTOR

U-2319 KYPROLIS IS INDICATED IN COMBINATION WITH DEXAMETHASONE OR WITH LENALIDOMIDE PLUS DEXAMETHASONE FOR THE TREATMENT OF PATIENTS WITH RELAPSED OR REFRACTORY MULTIPLE MYELOMA WHO HAVE RECEIVED ONE TO THREE LINES OF THERAPY

U-2320 KYPROLIS IS INDICATED AS A SINGLE AGENT FOR THE TREATMENT OF PATIENTS WITH RELAPSED OR REFRACTORY MULTIPLE MYELOMA WHO HAVE RECEIVED ONE OR MORE LINES OF THERAPY

U-2321 A METHOD OF APPLYING TRYPAN BLUE ONTO AN OUTER SURFACE OF THE ANTERIOR LENS CAPSULE TO FACILITATE REMOVAL OF THE LENS SUBSTANCE

U-2322 TREATMENT OF ADULT PATIENTS WITH MODERATELY TO SEVERELY ACTIVE ULCERATIVE COLITIS (UC)

U-2323 TREATMENT OF PATIENTS WITH CHRONIC LYMPHOCYTIC LEUKEMIA (CLL) OR SMALL LYMPHOCYTIC LYMPHOMA (SLL), WITH OR WITHOUT 17P DELETION, WHO HAVE RECEIVED AT LEAST ONE PRIOR THERAPY

PATENT AND EXCLUSIVITY TERMS

ADB 151 of 225

PATENT USE

U-2324 FOR SECONDARY PREVENTION OF CARDIOVASCULAR AND CEREBROVASCULAR EVENTS IN PATIENTS AT RISK OF DEVELOPING ASPIRIN-ASSOCIATED GASTRIC ULCERS

U-2325 EMERGENCY TREATMENT OF ALLERGIC REACTIONS (TYPE 1), INCLUDING ANAPHYLAXIS; A METHOD OF TREATING ALLERGIC REACTION, ANAPHYLAXIS, ANAPHYLACTIC SHOCK, OR COMBINATION THEREOF BY AN INJECTION OF AT LEAST ONE DOSAGE OF THE INJECTABLE LIQUID PHARMACEUTICAL

U-2326 TREATMENT OF NOCTURIA DUE TO NOCTURNAL POLYURIA IN ADULTS

U-2327 TREATMENT OF NOCTURIA DUE TO NOCTURNAL POLYURIA IN ADULTS, COMPRISING MONITORING A PATIENT'S SERUM SODIUM CONCENTRATION

U-2328 METHOD OF USING PLAZOMICIN TO TREAT BACTERIAL INFECTIONS

U-2329 METHOD OF ADMINISTERING A LOCAL ANESTHETIC PRIOR TO PERFORMING A DIAGNOSTIC OR SURGICAL PROCEDURE ON A SUBJECT WITH HEPATIC OR RENAL IMPAIRMENT

U-2330 METHOD OF TREATING MELANOMA

U-2331 INDICATED IN COMBINATION WITH ENCORAFENIB FOR THE TREATMENT OF MELANOMA

U-2332 INDICATED IN COMBINATION WITH ENCORAFENIB FOR THE TREATMENT OF MELANOMA MEDIATED BY A B-RAF PROTEIN KINASE

U-2333 INDICATED IN COMBINATION WITH ENCORAFENIB FOR THE TREATMENT OF MELANOMA WITH A BRAF MUTATION

U-2334 TREATMENT OF MELANOMA WITH A BRAF MUTATION

U-2335 TREATMENT OF MELANOMA

U-2336 TREATMENT OF MELANOMA MEDIATED BY A B-RAF PROTEIN KINASE

U-2337 INDICATED IN COMBINATION WITH BINIMETINIB FOR THE TREATMENT OF MELANOMA WITH A BRAF MUTATION

U-2338 MAINTAINING MEAN ARTERIAL PRESSURE OF ABOUT 65 MMHG OR ABOVE WITH ABOUT 1 NG/KG/MIN TO ABOUT 40 NG/KG/MIN ANGIOTENSIN II IN HYPOTENSIVE PATIENTS TREATED WITH VASOPRESSIN OR A VASOPRESSIN ANALOGUE AND REDUCING VASOPRESSIN OR VASOPRESSIN ANALOGUE USE

U-2339 USE OF A PHARMACEUTICAL COMPOSITION COMPRISING LINAGLIPTIN, METFORMIN AND A BASIC AMINO ACID TO TREAT TYPE 2 DIABETES MELLITUS

U-2340 TREATMENT OF POSTOPERATIVE INFLAMMATION

U-2341 METHOD OF RECONSTITUTING A LYOPHILIZED LIPOSOMAL COMPOSITION FOR ADMINISTERING CYTARABINE AND DAUNORUBICIN TO TREAT ADULTS WITH NEWLY-DIAGNOSED THERAPY-RELATED ACUTE MYELOID LEUKEMIA (T-AML) OR AML WITH MYELODYSPLASIA-RELATED CHANGES (AML-MRC)

U-2342 METHOD OF ADMINISTERING A RECONSTITUTED LIPOSOMAL COMPOSITION CONTAINING CYTARABINE AND DAUNORUBICIN TO TREAT ADULTS WITH NEWLY-DIAGNOSED THERAPY-RELATED ACUTE MYELOID LEUKEMIA (T-AML) OR AML WITH MYELODYSPLASIA-RELATED CHANGES (AML-MRC)

U-2343 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 12 YEARS AND OLDER, WHO ARE HOMOZYGOUS FOR THE F508DEL MUTATION OR HETEROZYGOUS FOR F508DEL AND A SECOND CFTR MUTATION PREDICTED TO BE RESPONSIVE TO TEZACAFTOR/IVACAFTOR, WITH TEZACAFTOR AND IVACAFTOR

U-2344 TREATMENT OF THROMBOCYTOPENIA IN ADULT PATIENTS WITH CHRONIC LIVER DISEASE WHO ARE SCHEDULED TO UNDERGO A PROCEDURE

U-2345 TREATMENT OF PATIENTS WITH CASTRATION-RESISTANT PROSTATE CANCER (CRPC)

U-2346 TREATMENT OF HUMAN SMALLPOX DISEASE CAUSED BY VARIOLA VIRUS IN ADULTS AND PEDIATRIC PATIENTS WEIGHING AT LEAST 13 KG

U-2347 TREATMENT OF TYPE 2 DIABETES MELLITUS IN A PATIENT WITH RENAL IMPAIRMENT AND FOR WHOM METFORMIN THERAPY IS INAPPROPRIATE BY ADMINISTERING LINAGLIPTIN WITHOUT DOSE ADJUSTMENT

U-2348 A METHOD FOR PREVENTION OF PREGNANCY

U-2349 FOR ONCE-DAILY MAINTENANCE TREATMENT OF ASTHMA AS PROPHYLACTIC THERAPY IN PATIENTS AGED 5 YEARS AND OLDER

U-2350 A METHOD OF TREATING A CANCER CHARACTERIZED BY AN IDH1 MUTATION WHERE THE CANCER IS ACUTE MYELOGENOUS LEUKEMIA (AML)

U-2351 TREATMENT OF ACUTE MYELOID LEUKEMIA (AML) WITH AN IDH1 MUTATION

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

- U-2352 TREATMENT OF HIV-1 INFECTION IN ADULTS WHO HAVE NO PRIOR ANTIRETROVIRAL TREATMENT HISTORY OR ARE VIROLOGICALLY SUPPRESSED ON A STABLE ANTIRETROVIRAL REGIMEN FOR AT LEAST 6 MONTHS
- U-2353 TX OF HIV-1 INFECTION USING A COMPOSITION CONTAINING A PK ENHANCER THAT INHIBITS CYTOCHROME P450 MONOOXYGENASES IN ADULTS WHO HAVE NO PRIOR ANTIRETROVIRAL TX HISTORY OR ARE VIROLOGICALLY SUPPRESSED ON A STABLE ANTIRETROVIRAL REGIMEN FOR AT LEAST 6 MONTHS
- U-2354 COMBINATION WITH OTHER ANTIRETROVIRALS (ATV) FOR TREATMENT OF HIV-1 IN ATV TREATMENT-EXPERIENCED PATIENTS 2 YEARS AND OLDER WITH EVIDENCE OF VIRAL REPLICATION AND HIV-1 STRAINS RESISTANT TO NON-NUCLEOSIDE REVERSE TRANSCRIPTASE INHIBITOR AND OTHER ATV
- U-2355 IN COMBINATION WITH AN AROMATASE INHIBITOR FOR THE TREATMENT OF PRE/PERIMENOPAUSAL OR POSTMENOPAUSAL WOMEN WITH HR-POSITIVE, HER2-NEGATIVE ADVANCED OR METASTATIC BREAST CANCER, AS INITIAL ENDOCRINE-BASED THERAPY
- U-2356 IN COMBINATION WITH FULVESTRANT FOR THE TREATMENT OF POSTMENOPAUSAL WOMEN WITH HR-POSITIVE, HER2-NEGATIVE ADVANCED OR METASTATIC BREAST CANCER, AS INITIAL ENDOCRINE BASED THERAPY OR FOLLOWING DISEASE PROGRESSION ON ENDOCRINE THERAPY
- U-2357 METHOD OF TREATING ATTENTION DEFICIT HYPERACTIVITY DISORDER (ADHD)
- U-2358 TREATMENT OF PATIENTS WITH HORMONE RECEPTOR (HR)-NEGATIVE BREAST CANCER WITH DELETERIOUS OR SUSPECTED DELETERIOUS GBRAM, HER2-NEGATIVE METASTATIC BREAST CANCER, WHO HAVE BEEN TREATED WITH CHEMOTHERAPY IN NEOADJUVANT, ADJUVANT, OR METASTATIC SETTING
- U-2359 TREATMENT OF PATIENTS WITH HORMONE RECEPTOR (HR)-POSITIVE BREAST CANCER WHO SHOULD HAVE BEEN TREATED WITH PRIOR ENDOCRINE THERAPY OR BE CONSIDERED INAPPROPRIATE FOR ENDOCRINE THERAPY
- U-2360 MANAGEMENT OF MODERATE TO SEVERE PAIN ASSOCIATED WITH ENDOMETRIOSIS
- U-2361 METHOD OF ADMINISTERING A GRANULATE FORMULATION OF 5-METHYL-1-PHENYL-2-(1H)-PYRIDONE AS RECITED IN CLAIM 1, TO TREAT IDIOPATHIC PULMONARY FIBROSIS
- U-2362 TREATMENT OF HEPATITIS C VIRUS (HCV) GENOTYPE 1, 2, 3, 4, 5, OR 6
- U-2363 ADMINISTRATION OF RISPERIDONE
- U-2364 TREATMENT OF HIV-1 INFECTION USING A COMPOSITION CONTAINING A PHARMACOKINETIC ENHANCER THAT INHIBITS CYTOCHROME P450 MONOOXYGENASE IN ADULTS WHO HAVE NO PRIOR ANTIRETROVIRAL TREATMENT HISTORY
- U-2365 TREATMENT OF HIV-1 INFECTION USING A COMPOSITION CONTAINING A PHARMACOKINETIC ENHANCER THAT INHIBITS CYTOCHROME P450 MONOOXYGENASE IN ADULTS WHO ARE VIROLOGICALLY SUPPRESSED ON A STABLE ANTIRETROVIRAL REGIMEN FOR AT LEAST 6 MONTHS
- U-2366 TREATMENT OF LIVER DISEASE THROUGH NUTRITION FOR PATIENTS UNDER THE AGE OF 12
- U-2367 USE FOR PATIENTS WITH PARENTERAL NUTRITION ASSOCIATED CHOLESTASIS OR PARENTERAL NUTRITION ASSOCIATED LIVER DISEASE
- U-2368 TOPICAL TREATMENT OF ACNE VULGARIS IN PATIENTS 9 YEARS OF AGE AND OLDER
- U-2369 FOR THE TREATMENT OF GENOTYPE 1, 4, 5 OR 6 CHRONIC HEPATITIS C VIRUS (HCV) INFECTION
- U-2370 FOR TREATMENT-NAIVE GENOTYPE 1 PATIENTS WITH CHRONIC HEPATITIS C VIRUS (HCV) INFECTION FOR A DURATION OF 8-WEEKS
- U-2371 THE TREATMENT OF FABRY PATIENTS
- U-2372 A METHOD OF REDUCING LEFT VENTRICULAR MASS INDEX (LVMI) IN A FABRY PATIENT BY ADMINISTERING MIGALASTAT
- U-2373 A METHOD OF REDUCING PODOCYTE GLOBOTRIAOSYL CERAMIDE (GL-3) IN A FABRY PATIENT BY ADMINISTERING MIGALASTAT
- U-2374 TREATMENT OF CYSTIC FIBROSIS IN A PATIENT AGE 2-5 YEARS WHO IS HOMOZYGOUS FOR THE F508DEL MUTATION IN THE CFTR GENE USING LUMACAFOR AND IVACAFOR
- U-2375 TREATMENT OF CYSTIC FIBROSIS IN A PATIENT AGE 2-5 YEARS WHO IS HOMOZYGOUS FOR THE F508DEL MUTATION IN THE CFTR GENE USING LUMACAFOR FORM I AND IVACAFOR
- U-2376 TREATMENT OF CYSTIC FIBROSIS IN A PATIENT AGE 2-5 YEARS WHO IS HOMOZYGOUS FOR THE F508DEL MUTATION IN THE CFTR GENE USING LUMACAFOR AND A SOLID COMPOSITION COMPRISING AMORPHOUS AND LESS THAN ABOUT 30% CRYSTALLINE IVACAFOR
- U-2377 USE OF VITAL DYE FOR FACILITATING SURGICAL PROCEDURES FOR VITREO-RETINAL SURGERY

PATENT AND EXCLUSIVITY TERMS

ADB 153 of 225

PATENT USE

U-2378 TREATMENT OF POLYNEUROPATHY OF HEREDITARY TRANSTHYRETIN-MEDIATED AMYLOIDOSIS

U-2379 USE IN IDENTIFICATION OF INTRAOCULAR MEMBRANES TO FACILITATE REMOVAL DURING OPHTHALMIC SURGERY

U-2380 TREATMENT OF COMPLICATED INTRA-ABDOMINAL INFECTIONS IN PATIENTS 18 YEARS OF AGE AND OLDER

U-2381 TREATMENT IN COMBINATION WITH A GNRH AGONIST OF NON-METASTATIC, CASTRATION-RESISTANT PROSTATE CANCER (NM-CRPC)

U-2382 TREATMENT IN COMBINATION WITH A GNRH AGONIST OF HIGH RISK NON-METASTATIC, CASTRATION-RESISTANT PROSTATE CANCER (NM-CRPC)

U-2383 METHOD OF CONTROLLING GLYCEMIA IN A DIABETIC PATIENT WITH DELAYED OR PROLONGED FOOD ABSORPTION BY ADMINISTERING 50 TO 75% OF A PREDETERMINED DOSE OF INSULIN-FDKP AT MEALTIME, AND ADMINISTERING REMAINDER OF DOSE 30-120 MINUTES AFTER BEGINNING OF MEAL

U-2384 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING A DOSAGE FORM COMPRISING MICRONIZED BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 1 AND 10

U-2385 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING A DOSAGE FORM COMPRISING MICRONIZED BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 1,10 AND 11

U-2386 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING A DOSAGE FORM COMPRISING BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 12 AND 19

U-2387 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING A DOSAGE FORM COMPRISING BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 12, 19 AND 20

U-2388 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING A DOSAGE FORM COMPRISING BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 21 AND 28

U-2389 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING A DOSAGE FORM COMPRISING BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 21, 28, AND 29

U-2390 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING A DOSAGE FORM COMPRISING BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 30 AND 41

U-2391 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING A DOSAGE FORM COMPRISING BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 30, 41, AND 42

U-2392 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING A DOSAGE FORM COMPRISING BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 43 AND 50

U-2393 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING A DOSAGE FORM COMPRISING BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 43, 50 AND 51

U-2394 FOR USE IN COMBINATION WITH OTHER ANTIRETROVIRAL AGENTS FOR THE TREATMENT OF HIV-1 INFECTION IN ADULT PATIENTS WITH NO PRIOR ANTIRETROVIRAL TREATMENT HISTORY

U-2395 FOR THE TREATMENT OF HIV-1 INFECTION IN ADULT PATIENTS WITH NO PRIOR ANTIRETROVIRAL TREATMENT HISTORY

U-2396 TREATMENT OF CYSTIC FIBROSIS IN A PATIENT AGE 2-5 YEARS WHO IS HOMOZYGOUS FOR THE F508DEL MUTATION IN THE CFTR GENE USING THE DOSAGE UNIT COMPRISING LUMACAFTOR AS RECITED IN CLAIM 1 OF US PATENT 8716338 AND IVACAFTOR

U-2397 TREATMENT OF CYSTIC FIBROSIS IN A PATIENT AGE 2-5 YEARS WHO IS HOMOZYGOUS FOR THE F508DEL MUTATION IN THE CFTR GENE USING THE DOSAGE UNIT COMPRISING LUMACAFTOR AND IVACAFTOR AS RECITED IN CLAIM 1 OF US PATENT 9192606

U-2398 TOPICAL TREATMENT OF PRIMARY AXILLARY HYPERHIDROSIS IN ADULTS AND PEDIATRIC PATIENTS 9 YEARS OF AGE AND OLDER

U-2399 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS 12 YEARS AND OLDER, WITH A F508DEL OR G551D CFTR GENE MUTATION AND A A455E, 2789+5G->A, OR 3849+10KBC->T MUTATION, COMPRISING CONCURRENT COADMINISTRATION OF THE COMPOSITIONS OF CLAIM 1 OF U.S. PATENT 10058546

PATENT AND EXCLUSIVITY TERMS

ADB 154 of 225

PATENT USE

U-2400 REDUCING ELEVATED INTRAOCULAR PRESSURE IN PATIENTS WITH OPEN-ANGLE GLAUCOMA OR OCULAR HYPERTENSION

U-2401 A METHOD OF TREATING AMYOTROPHIC LATERAL SCLEROSIS IN A PATIENT IN NEED OF SUCH TREATMENT, SAID METHOD COMPRISING ADMINISTERING TO SAID PATIENT AN EFFECTIVE AMOUNT OF A SUSPENSION ACCORDING TO CLAIM 1

U-2402 TREATMENT OF SCHIZOPHRENIA BY RAPID AND CONTINUOUS INTRAMUSCULAR INJECTION

U-2403 TREATMENT OF PSORIASIS USING A DOSAGE TITRATION SCHEDULE

U-2404 METHOD OF DELIVERING SUMATRIPTAN TO A NASAL CAVITY

U-2405 A METHOD FOR TREATING A BACTERIAL INFECTION IN INFLAMMATORY LESIONS OF NON-NODULAR MODERATE TO SEVERE ACNE VULGARIS PATIENTS 9 YEARS OF AGE AND OLDER COMPRISING ADMINISTERING AN EFFECTIVE AMOUNT OF SARECYCLINE HYDROCHLORIDE

U-2406 A METHOD FOR TREATING A PATIENT 9 YEARS OF AGE AND OLDER SUFFERING FROM AN INFLAMMATORY SKIN DISORDER OF NON-NODULAR MODERATE TO SEVERE ACNE VULGARIS COMPRISING ADMINISTERING AN EFFECTIVE AMOUNT OF SARECYCLINE HYDROCHLORIDE

U-2407 A METHOD FOR TREATING ACNE IN INFLAMMATORY LESIONS OF NON-NODULAR MODERATE TO SEVERE ACNE VULGARIS PATIENTS 9 YEARS OF AGE AND OLDER COMPRISING ADMINISTERING AN EFFECTIVE AMOUNT OF SARECYCLINE HYDROCHLORIDE CRYSTALLINE SALT

U-2408 A METHOD FOR TREATING A BACTERIAL INFECTION IN INFLAMMATORY LESIONS OF NON-NODULAR MODERATE TO SEVERE ACNE VULGARIS PATIENTS 9 YEARS OF AGE AND OLDER COMPRISING ADMINISTERING AN EFFECTIVE AMOUNT OF SARECYCLINE HYDROCHLORIDE CRYSTALLINE SALT

U-2409 A METHOD FOR TREATING ACNE IN INFLAMMATORY LESIONS OF NON-NODULAR MODERATE TO SEVERE ACNE VULGARIS PATIENTS 9 YEARS OF AGE AND OLDER COMPRISING ADMINISTERING SARECYCLINE HYDROCHLORIDE IN 60 MG, 100 MG OR 150 MG EQUIVALENT DOSES

U-2410 TREATMENT OF ADULT PATIENTS FOR WHOM TREATMENT WITH BOTH AMLODIPINE FOR HYPERTENSION AND CELECOXIB FOR OSTEOARTHRITIS ARE APPROPRIATE

U-2411 TREATMENT OF CYSTIC FIBROSIS IN A PATIENT AGE 12 YEARS OR OLDER WHO IS HOMOZYGOUS FOR THE F508DEL MUTATION IN THE CFTR GENE USING THE TABLET COMPRISING LUMACAFTOR AS RECITED IN CLAIM 1, 19, OR 21 OF U.S. PATENT NO. 10,076,513 AND IVACAFTOR

U-2412 FOR THE TREATMENT OF PATIENTS WITH CHRONIC LYMPHOCYTIC LEUKEMIA (CLL) AND/OR SMALL LYMPHOCYTIC LEUKEMIA (SLL)

U-2413 FOR THE TREATMENT OF PATIENTS WITH FOLLICULAR LYMPHOMA (FL)

U-2414 TREATING MYCOBACTERIUM AVIUM COMPLEX (MAC) LUNG DISEASE IN ADULTS AS PART OF A COMBINATION DRUG REGIMEN

U-2415 TREATING MYCOBACTERIUM AVIUM COMPLEX (MAC) LUNG DISEASE IN ADULTS AS PART OF A COMBINATION ANTIBACTERIAL DRUG REGIMEN

U-2416 TREATING MYCOBACTERIUM AVIUM COMPLEX (MAC) LUNG DISEASE IN ADULTS WITH CYSTIC FIBROSIS AS PART OF A COMBINATION DRUG REGIMEN

U-2417 TREATING MYCOBACTERIUM AVIUM COMPLEX (MAC) LUNG DISEASE IN NON-CYSTIC FIBROSIS ADULTS AS PART OF A COMBINATION ANTIBACTERIAL DRUG REGIMEN

U-2418 METHOD OF ADMINISTERING TESTOSTERONE ENANTHATE SUBCUTANEOUSLY

U-2419 METHOD OF OPERATING AN INJECTION DEVICE

U-2420 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGES 12 AND OLDER, WHO HAVE TWO COPIES OF THE F508DEL MUTATION OR ONE F508DEL MUTATION AND A CFTR MUTATION PREDICTED TO BE RESPONSIVE TO TEZACAFTOR/IVACAFTOR, WITH THE COMPOSITION OF CLAIM 1 OF US 10,081,621

U-2421 USE IN COMBINATION WITH CLOBAZAM FOR THE TREATMENT OF SEIZURES IN PATIENTS WITH DRAVET SYNDROME

U-2422 USE IN COMBINATION WITH CLOBAZAM FOR THE TREATMENT OF SEIZURES IN PATIENTS WITH LENNOX GASTAUT SYNDROME WHO HAVE BEEN PREVIOUSLY TREATED WITH CLOBAZAM

U-2423 USE IN COMBINATION WITH CLOBAZAM FOR THE TREATMENT OF SEIZURES IN PATIENTS WITH DRAVET SYNDROME WHO HAVE BEEN PREVIOUSLY TREATED WITH CLOBAZAM

U-2424 USE IN COMBINATION WITH CLOBAZAM FOR TREATMENT OF SEIZURES IN PATIENTS WITH LENNOX GASTAUT SYNDROME

U-2425 USE FOR THE TREATMENT OF CONVULSIVE SEIZURES IN PATIENTS WITH DRAVET SYNDROME

U-2426 USE FOR THE TREATMENT OF CONVULSIVE SEIZURES IN PATIENTS WITH LENNOX GASTAUT SYNDROME

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-2427 USE FOR THE TREATMENT OF DROP SEIZURES IN PATIENTS WITH DRAVET SYNDROME

U-2428 TREATMENT OF PARTIAL-ONSET SEIZURES WITH OR WITHOUT SECONDARILY GENERALIZED SEIZURES IN PATIENTS WITH EPILEPSY 4 YEARS OF AGE AND OLDER

U-2429 TREATMENT OF PRIMARY GENERALIZED TONIC-CLONIC SEIZURES AS ADJUNCTIVE THERAPY IN PATIENTS WITH EPILEPSY 12 YEARS OF AGE AND OLDER

U-2430 TREATMENT OF POLYNEUROPATHY OF HEREDITARY TRANSTHYRETIN AMYLOIDOSIS

U-2431 TREATMENT OF CHRONIC OBSTRUCTIVE PULMONARY DISEASE (COPD), INCLUDING CHRONIC BRONCHITIS

U-2432 LONG-TERM, MAINTENANCE TREATMENT OF BRONCHOSPASM ASSOCIATED WITH CHRONIC OBSTRUCTIVE PULMONARY DISEASE (COPD)

U-2433 METHOD OF TREATING A BIOLOGICAL RHYTHM DISORDER, SUCH AS INSOMNIA

U-2434 USE IN COMBINATION WITH LENALIDOMIDE AND DEXAMETHASONE FOR THE TREATMENT OF PATIENTS WITH MULTIPLE MYELOMA WHO HAVE RECEIVED AT LEAST ONE PRIOR THERAPY

U-2435 REDUCTION OF RISK OF MAJOR CARDIOVASCULAR EVENTS (CV DEATH, MI, AND STROKE) IN CHRONIC CAD OR PAD

U-2436 USE IN THE TREATMENT OF MAJOR DEPRESSIVE DISORDER TO IMPROVE TREATMENT EMERGENT SEXUAL DYSFUNCTION (TESD) INDUCED BY PRIOR SEROTONIN REUPTAKE INHIBITOR TREATMENT

U-2437 TREATMENT OF ADULT PATIENTS WITH DELETERIOUS OR SUSPECTED DELETERIOUS GERMLINE BREAST CANCER SUSCEPTIBILITY GENE (BRCA)-MUTATED (GBRCAM) HUMAN EPIDERMAL GROWTH FACTOR RECEPTOR 2 (HER2)-NEGATIVE LOCALLY ADVANCED OR METASTATIC BREAST CANCER

U-2438 CARDIOVASCULAR OUTCOMES TRIAL OF LIRAGLUTIDE 1.8 MG IN PATIENTS WITH TYPE 2 DIABETES AND CARDIOVASCULAR DISEASE

U-2439 TREATMENT OF MENOPAUSE SYMPTOMS, INCLUDING VASOMOTOR SYMPTOMS

U-2440 FOR THE MAINTENANCE TREATMENT OF PATIENTS WITH CHRONIC OBSTRUCTIVE PULMONARY DISEASE (COPD)

U-2441 REDUCTION OF RISK OF MAJOR ADVERSE CARDIOVASCULAR EVENTS IN THE TREATMENT OF TYPE 2 DIABETES MELLITUS PATIENTS

U-2442 USE FOR THE TREATMENT OF ATONIC SEIZURES IN PATIENTS WITH LENNOX-GASTAUT SYNDROME

U-2443 USE FOR THE TREATMENT OF ATONIC SEIZURES IN PATIENTS WITH DRAVET SYNDROME

U-2444 TREATMENT OF SUBJECTS HAVING BACTERIAL SKIN OR SKIN STRUCTURE INFECTION

U-2445 TREATMENT IN COMBINATION WITH AZACITIDINE OR DECITABINE OR LOW-DOSE CYTARABINE OF NEWLY-DIAGNOSED ACUTE MYELOID LEUKEMIA (AML) IN ADULTS WHO ARE AGE 75 YEARS OR OLDER, OR WHO HAVE COMORBIDITIES THAT PRECLUDE USE OF INTENSIVE INDUCTION CHEMOTHERAPY

U-2446 TREATMENT OF ADULT PATIENTS WITH CHRONIC LYMPHOCYTIC LEUKEMIA (CLL) OR SMALL LYMPHOCYTIC LYMPHOMA (SLL), WITH OR WITHOUT 17P DELETION, WHO HAVE RECEIVED AT LEAST ONE PRIOR THERAPY

U-2447 TREATMENT OF SEVERE HYPERTRIGLYCERIDEMIA (500 MG/DL) IN ADULT PATIENTS AS AN ADJUNCT TO DIET

U-2448 TREATMENT OF TRAVELERS' DIARRHEA CAUSED BY NON-INVASIVE STRAINS OF ESCHERICHIA COLI IN ADULTS

U-2449 TREATMENT OF BACTERIAL SKIN AND SKIN STRUCTURE INFECTION

U-2450 POSITRON EMISSION TOMOGRAPHY DIAGNOSTIC AGENT IN ADULTS WITH SUSPECTED PROSTATE CANCER RECURRENCE BASED ON ELEVATED BLOOD PROSTATE SPECIFIC ANTIGEN LEVELS FOLLOWING PRIOR TREATMENT

U-2451 TREATMENT OF THROMBOCYTOPENIA IN ADULT AND PEDIATRIC PATIENTS 1 YEAR AND OLDER WITH CHRONIC IMMUNE (IDIOPATHIC) THROMBOCYTOPENIA (ITP)

U-2452 COMBINATION WITH IMMUNOSUPPRESSIVE THERAPY FOR FIRST-LINE TREATMENT OF ADULT AND PEDIATRIC PATIENTS 2 YEARS AND OLDER WITH SEVERE APLASTIC ANEMIA

U-2453 TREATMENT OF FUNGAL INFECTIONS, INCLUDING BLASTOMYCOSIS, HISTOPLASMOSIS, AND ASPERGILLOSIS

U-2454 USE FOR THE TREATMENT OF DROP SEIZURES IN PATIENTS WITH LENNOX-GASTAUT SYNDROME

U-2455 USE IN COMBINATION WITH CLOBAZAM FOR TREATMENT OF DROP SEIZURES IN PATIENTS WITH LENNOX GASTAUT SYNDROME

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-2456 TREATMENT OF ACUTE MYELOID LEUKEMIA (AML)

U-2457 REINITIATION OF SCHIZOPHRENIA TREATMENT FOLLOWING A MISSED DOSE MORE THAN 9 MONTHS AGO

U-2458 REINITIATION OF SCHIZOPHRENIA TREATMENT FOLLOWING A MISSED DOSE 4-9 MONTHS AGO

U-2459 TREATMENT OF DYSKINESIA AND DECREASING OFF TIME IN PATIENTS WITH PARKINSON'S DISEASE RECEIVING LEVODOPA-BASED THERAPY, WITH OR WITHOUT CONCOMITANT DOPAMINERGIC MEDICATIONS

U-2460 VISUALIZATION OF VESSELS, BLOOD FLOW AND TISSUE PERFUSION OF CORONARY ARTERY BYPASS GRAFT IN VASCULAR, GASTROINTESTINAL, ORGAN TRANSPLANT, AND PLASTIC, MICRO- AND RECONSTRUCTIVE, INCLUDING MINIMALLY INVASIVE, SURGERY

U-2461 VISUALIZATION OF VESSELS, BLOOD FLOW AND TISSUE PERFUSION OF CARDIOVASCULAR BYPASS GRAFT AND VASCULATURE IN VASCULAR, GASTROINTESTINAL, ORGAN TRANSPLANT, AND PLASTIC, MICRO- AND RECONSTRUCTIVE, INCLUDING MINIMALLY INVASIVE, SURGERY

U-2462 VISUALIZATION OF VESSELS, BLOOD FLOW AND TISSUE PERFUSION OF VESSEL WITH ARTERIOVENOUS MALFORMATION IN VASCULAR, GASTROINTESTINAL, ORGAN TRANSPLANT, AND PLASTIC, MICRO- AND RECONSTRUCTIVE, INCLUDING MINIMALLY INVASIVE, SURGERY

U-2463 VISUALIZATION OF VESSELS, BLOOD FLOW AND TISSUE PERFUSION IN SURGICAL FLAPS IN VASCULAR, GASTROINTESTINAL, ORGAN TRANSPLANT, AND PLASTIC, MICRO- AND RECONSTRUCTIVE, INCLUDING MINIMALLY INVASIVE, SURGERY

U-2464 VISUALIZATION OF VESSELS, BLOOD FLOW AND TISSUE PERFUSION OF TRANSPLANTED ORGAN OR ATTACHED VESSEL IN VASCULAR, GASTROINTESTINAL, ORGAN TRANSPLANT, AND PLASTIC, MICRO- AND RECONSTRUCTIVE, INCLUDING MINIMALLY INVASIVE, SURGERY

U-2465 VISUALIZATION OF VESSELS, BLOOD FLOW AND TISSUE PERFUSION OF VESSEL GRAFT IN VASCULAR, GASTROINTESTINAL, ORGAN TRANSPLANT, AND PLASTIC, MICRO- AND RECONSTRUCTIVE, INCLUDING MINIMALLY INVASIVE, SURGERY

U-2466 VISUALIZATION OF VESSELS, BLOOD FLOW AND TISSUE PERFUSION OF DONOR ORGAN OR ATTACHED VESSEL IN VASCULAR, GASTROINTESTINAL, ORGAN TRANSPLANT, AND PLASTIC, MICRO- AND RECONSTRUCTIVE, INCLUDING MINIMALLY INVASIVE, SURGERY

U-2467 VISUALIZATION OF EXTRAHEPATIC BILIARY DUCT ATTACHED TO DONOR ORGAN IN PATIENTS 12 YEARS AND OLDER

U-2468 VISUALIZATION OF EXTRAHEPATIC BILIARY DUCT ATTACHED TO TRANSPLANTED ORGAN IN PATIENTS 12 YEARS AND OLDER

U-2469 METHOD OF TREATING CANCEROUS SOLID TUMORS

U-2470 METHOD OF TREATING SOLID TUMORS THAT EXHIBIT AN NTRK GENE FUSION

U-2471 METHOD OF TREATING SOLID TUMORS THAT EXHIBIT AN NTRK FUSION GENE IN A PEDIATRIC PATIENT

U-2472 METHOD OF TREATING NEUROBLASTOMA, GLIOMA, THYROID, AND BREAST CANCER SOLID TUMORS THAT EXHIBIT AN NTRK GENE FUSION

U-2473 METHOD OF TREATING CMN, IFS, HGG, DIPGS, PTC, SOFT TISSUE SARCOMA, AND SPINDLE CELL SARCOMA SOLID TUMORS EXHIBITING AN NTRK GENE FUSION IN A PEDIATRIC PATIENT WITH AN ORAL SOLUTION

U-2474 METHOD OF TREATING SOLID TUMORS THAT EXHIBIT AN NTRK GENE FUSION AFTER SURGICAL RESECTION

U-2475 METHOD OF TREATING SOLID TUMORS THAT EXHIBIT AN NTRK GENE FUSION IN A PEDIATRIC PATIENT

U-2476 USE OF A DELIVERY DEVICE TO DELIVER A DOSE OF NALOXONE

U-2477 TREATMENT OF NON-24 HOUR SLEEP-WAKE DISORDER BY AVOIDING THE USE OF TASIMELTEON IN COMBINATION WITH CYP1A2 STRONG INHIBITORS

U-2478 METHOD FOR THE INDUCTION OF LOCAL ANESTHESIA PRIOR TO PERFORMING A PROCEDURE ON, THROUGH, OR ADJACENT TO THE MUCOUS MEMBRANES

U-2479 METHOD OF ADMINISTERING A LOCAL ANESTHETIC TO THE MUCOUS MEMBRANES

U-2480 MAINTENANCE TREATMENT OF GBRCA- OR SBRCA-MUTATED ADVANCED EPITHELIAL OVARIAN, FALLOPIAN TUBE OR PRIMARY PERITONEAL CANCER WHO ARE IN A COMPLETE OR PARTIAL RESPONSE TO FIRST-LINE PLATINUM-BASED CHEMOTHERAPY

U-2481 TREATMENT OF DELETERIOUS OR SUSPECTED DELETERIOUS GERMLINE BRCA-MUTATED ADVANCED OVARIAN CANCER WHO HAVE BEEN TREATED WITH THREE OR MORE PRIOR LINES OF CHEMOTHERAPY

PATENT AND EXCLUSIVITY TERMS

ADB 157 of 225

PATENT USE

- U-2482 TREATMENT OF HR-NEGATIVE, HER-2 NEGATIVE, GBRCA-MUTATED METASTATIC BREAST CANCER, WHO HAVE BEEN TREATED WITH CHEMOTHERAPY IN THE NEOADJUVANT, ADJUVANT, OR METASTATIC SETTING
- U-2483 TREATMENT OF HR-POSITIVE, HER-2 NEGATIVE, GBRCA-MUTATED METASTATIC BREAST CANCER, WHO HAVE BEEN TREATED WITH CHEMOTHERAPY IN THE NEOADJUVANT, ADJUVANT, OR METASTATIC SETTING, AND WITH ENDOCRINE THERAPY OR ARE INAPPROPRIATE FOR ENDOCRINE THERAPY
- U-2484 INTERMITTENT TREATMENT OF OFF EPISODES IN PATIENTS WITH PARKINSON'S DISEASE TREATED WITH CARBIDOPA/LEVODOPA BY INHALATION OF LEVODOPA POWDER PARTICLES
- U-2485 INTERMITTENT TREATMENT OF OFF EPISODES IN PATIENTS WITH PARKINSON'S DISEASE TREATED WITH CARBIDOPA/LEVODOPA BY INHALATION OF LEVODOPA POWDER PARTICLES THROUGH A SINGLE BREATH ACTIVATED STEP
- U-2486 INTERMITTENT TREATMENT OF OFF EPISODES IN PATIENTS WITH PARKINSON'S DISEASE WITH A POWDER INHALER
- U-2487 DEXTENZA IS APPROVED FOR THE TREATMENT OF OCULAR PAIN FOLLOWING OPHTHALMIC SURGERY
- U-2488 TREATMENT OF PATIENTS WITH HEPATOCELLULAR CARCINOMA (HCC) WHO HAVE BEEN PREVIOUSLY TREATED WITH SORAFENIB
- U-2489 TREATMENT OF MODERATE TO SEVERE OPIOID USE DISORDER
- U-2490 TREATMENT OF COMPLICATED URINARY TRACT INFECTION (CUTI) INCLUDING PYELONEPHRITIS CAUSED BY THE FOLLOWING SUSCEPTIBLE MICROORGANISMS: ESCHERICHIA COLI, KLEBSIELLA PNEUMONIA, AND ENTEROBACTER CLOACAE SPECIES COMPLEX
- U-2491 A METHOD FOR DELIVERING A COMPOSITION TO A MUCUS MEMBRANE
- U-2492 A METHOD FOR DELIVERING A PHARMACEUTICAL AGENT ACROSS A MUCOSAL BARRIER
- U-2493 A METHOD FOR TREATING INFLAMMATION AND/OR OTHER DISORDERS IN AN EYE OF A PATIENT
- U-2494 INDICATED FOR THE TREATMENT OF VENTRICULAR ARRHYTHMIAS, SUCH AS SUSTAINED VENTRICULAR TACHYCARDIA, THAT IN THE JUDGEMENT OF THE PHYSICIAN ARE LIFE-THREATENING
- U-2495 VENTRICULAR FIBRILLATION
- U-2496 IMPROVEMENT IN GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS BY USE OF A PEN INJECTOR WITH A THREADED DRIVE SLEEVE
- U-2497 TREATMENT OF DRUG-INDUCED EXTRAPYRAMIDAL REACTION IN ADULT PATIENTS
- U-2498 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGES 12 AND OLDER, WHO HAVE TWO COPIES OF THE F508DEL MUTATION OR ONE F508DEL MUTATION AND A CFTR MUTATION PREDICTED TO BE RESPONSIVE TO TEZACAFTOR/IVACAFTOR, WITH THE COMPOSITION OF CLAIM 1 OF US 10,206,877
- U-2499 METHOD OF REDUCING ADVERSE EFFECTS IN PATIENTS SUFFERING FROM EXCESSIVE DAYTIME SLEEPINESS AND/OR CATAPLEXY IN NARCOLEPSY WHO ARE CONCOMITANTLY ADMINISTERED SODIUM OXYBATE AND DIVALPROEX SODIUM
- U-2500 USE OF A DELIVERY DEVICE TO DELIVER A BIOEQUIVALENT DOSE OF A NALOXONE COMPOSITION VIA A NEEDLE
- U-2501 TREATMENT OF PARTIAL-ONSET SEIZURES
- U-2502 TREATMENT OF TREATMENT-RESISTANT DEPRESSION IN ADULT IN CONJUNCTION WITH AN ORAL ANTIDEPRESSANT
- U-2503 TREATMENT OF ADULTS WITH METASTATIC GASTRIC OR GJA PREVIOUSLY TREATED WITH AT LEAST TWO PRIOR LINES OF CHEMOTHERAPY THAT INCLUDED A FLUOROPYRIMIDINE, A PLATINUM, EITHER A TAXANE OR IRINOTECAN, AND IF APPROPRIATE, HER2/NEU-TARGETED THERAPY
- U-2504 TREATMENT OF HR-POSITIVE, HER2-NEGATIVE ADVANCED OR METASTATIC BREAST CANCER IN POSTMENOPAUSAL WOMEN IN COMBINATION WITH RIBOCICLIB AS INITIAL ENDOCRINE BASED THERAPY OR FOLLOWING DISEASE PROGRESSION ON ENDOCRINE THERAPY
- U-2505 TREATMENT OF PRE/PERIMENOPAUSAL WOMEN WITH HORMONE RECEPTOR (HR)-POSITIVE, HUMAN EPIDERMAL GROWTH FACTOR RECEPTOR 2 (HER2)-NEGATIVE ADVANCED OR METASTATIC BREAST CANCER
- U-2506 METHOD OF TREATING TESTOSTERONE DEFICIENCY
- U-2507 METHOD OF TREATING ACUTE BACTERIAL SKIN AND SKIN STRUCTURE INFECTIONS (ABSSSI) CAUSED BY DESIGNATED SUSCEPTIBLE BACTERIA

PATENT AND EXCLUSIVITY TERMS

ADB 158 of 225

PATENT USE

- U-2508 A METHOD OF TREATING BACTERIAL INFECTIONS IN COMPLICATED INTRA-ABDOMINAL INFECTION AND COMPLICATED URINARY TRACT INFECTION, INCLUDING PYELONEPHRITIS, PATIENTS COMPRISING ADMINISTERING A BACTERICIDALLY EFFECTIVE AMOUNT OF AVIBACTAM SODIUM
- U-2509 A METHOD OF TREATING A BACTERIAL INFECTION IN COMPLICATED INTRA-ABDOMINAL INFECTION (CIAI) AND COMPLICATED URINARY TRACT INFECTION (CUTI), INCLUDING PYELONEPHRITIS, PATIENTS COMPRISING ADMINISTERING AN EFFECTIVE AMOUNT OF AVIBACTAM SODIUM
- U-2510 A METHOD FOR CONTRACEPTION COMPRISING THE STEP OF ORAL ADMINISTRATION A DOSAGE OF 20 MG TO 30 MG OF ULIPRISTAL ACETATE TO A WOMAN WITHIN 72 HOURS AND UP TO 120 HOURS AFTER AN UNPROTECTED INTERCOURSE
- U-2511 A METHOD OF TREATING MULTIPLE SCLEROSIS BY ADMINISTERING SIPONIMOD USING A TITRATION SCHEME TO REACH A MAINTENANCE DOSE
- U-2512 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 12 YEARS AND OLDER, WHO ARE HOMOZYGOUS FOR THE F508DEL MUTATION OR HAVE AT LEAST ONE CFTR MUTATION THAT IS RESPONSIVE TO TEZACAFTOR/IVACAFTOR, WITH AN EFFECTIVE AMOUNT OF TEZACAFTOR AND IVACAFTOR
- U-2513 MAINTENANCE TREATMENT OF CHRONIC OBSTRUCTIVE PULMONARY DISEASE (COPD)
- U-2514 MAINTENANCE TREATMENT OF BRONCHOSPASM ASSOCIATED WITH CHRONIC OBSTRUCTIVE PULMONARY DISEASE (COPD)
- U-2515 PALBOCICLIB FOR HR-POS. HER2-NEG. ADVANCED OR METASTATIC BREAST CANCER IN COMBO WITH AN AROMATASE INHIBITOR AS INITIAL ENDOCRINE-BASED THERAPY IN POSTMENOPAUSAL WOMEN OR MEN, OR WITH FULVESTRANT IN PTS WITH DISEASE PROGRESSION AFTER ENDOCRINE THERAPY
- U-2516 A METHOD FOR REDUCING SERUM GLUCOSE LEVELS IN ADULTS WITH TYPE 2 DIABETES MELLITUS
- U-2517 A METHOD FOR REDUCING SERUM GLUCOSE LEVELS IN ADULTS WITH TYPE 2 DIABETES MELLITUS
- U-2518 TREATMENT OF ADULTS WITH LOCALLY ADVANCED OR METASTATIC UROTHELIAL CARCINOMA THAT HAS SUSCEPTIBLE FGFR3 OR FGFR2 GENETIC ALTERATIONS AND PROGRESSED DURING OR FOLLOWING PRIOR PLATINUM-CONTAINING CHEMOTHERAPY
- U-2519 TREATMENT OF ADULT PATIENTS WITH MANTLE CELL LYMPHOMA WHO HAVE RECEIVED AT LEAST ONE PRIOR THERAPY BY ADMINISTERING 100 MG OF ACALABRUTINIB TWICE DAILY
- U-2520 TREATING MS WITH ORAL CLADRIBINE ACC. TO THE STEPS (I) INDUCTION PERIOD WITH ABOUT 1.7 MG/KG-3.5 MG/KG CLADRIBINE; (II) CLADRIBINE-FREE PERIOD OF ABOUT 8-10 MONTHS; (III) MAINTENANCE PERIOD WITH ABOUT 1.7 MG/KG CLADRIBINE; (IV) CLADRIBINE-FREE PERIOD
- U-2521 TREATMENT OF MS WITH A TABLET WITH AN ADMIXTURE OF (A) AN AMORPHOUS INCLUSION COMPLEX OF CLADRIBINE AND HYDROXYPROPYL-B-CYCLODEXTRIN AND (B) AMORPHOUS FREE CLADRIBINE AND CYCLODEXTRIN AS A NON-INCLUSION COMPLEX, CLADRIBINE/CYCLODEXTRIN 1:10-1:16 W/W
- U-2522 TREATING RRMS OR SPMS WITH ORAL CLADRIBINE: (I) 2-4 MONTHS INDUCTION WITH 1.7 MG/KG - 3.5 MG/KG CLADRIBINE; (II) CLADRIBINE-FREE PERIOD OF ABOUT 8-10 MONTHS; (III) 2-4 MONTHS MAINTENANCE WITH ABOUT 1.7 MG/KG CLADRIBINE; (IV) CLADRIBINE-FREE PERIOD
- U-2523 TREATMENT OF MS WITH AN ADMIXTURE OF (A) AN AMORPHOUS INCLUSION COMPLEX OF CLADRIBINE (2CDA) AND CYCLODEXTRIN AND (B) AMORPHOUS FREE 2CDA AND CYCLODEXTRIN AS A NON-INCLUSION COMPLEX, FORMULATED AS A SOLID ORAL FORM, W/O SIGN. AMOUNTS OF CRYST. 2CDA
- U-2524 TREATMENT OF THE CARDIOMYOPATHY OF WILD TYPE OR HEREDITARY TRANSTHYRETIN-MEDIATED AMYLOIDOSIS (ATTR-CM)
- U-2525 TREATMENT OF PATIENTS WITH AN OVERACTIVE BLADDER WITH SYMPTOMS OF URINARY FREQUENCY, URGENCY OR URGE INCONTINENCE
- U-2526 ACUTE TREATMENT OF INTERMITTENT, STEREOTYPIC EPISODES OF FREQUENT SEIZURE ACTIVITY (I.E., SEIZURE CLUSTERS, ACUTE REPETITIVE SEIZURES) THAT ARE DISTINCT FROM A PATIENT'S USUAL SEIZURE PATTERN IN PATIENTS WITH EPILEPSY 12 YEARS OF AGE AND OLDER
- U-2527 TREATMENT OF CYSTIC FIBROSIS USING IVACAFTOR IN A PATIENT AGE 6 MONTHS TO <6 YEARS WHO HAS ONE MUTATION IN THE CFTR GENE THAT IS RESPONSIVE TO IVACAFTOR BASED ON CLINICAL AND/OR IN VITRO ASSAY DATA

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

- U-2528 TREATMENT OF CYSTIC FIBROSIS USING IVACAFTOR IN A PATIENT AGE 6 MONTHS TO <6 YEARS WHO HAS A R117H MUTATION IN THE CFTR GENE
- U-2529 TREATMENT OF A MODERATE MILD CLINICAL PHENOTYPE OF CF USING IVACAFTOR IN A PATIENT AGE 6 MONTHS TO <6 YEARS WHO HAS ONE CFTR MUTATION RESPONSIVE TO IVACAFTOR BASED ON CLINICAL AND/OR IN VITRO ASSAY DATA
- U-2530 TREATMENT OF CF IN A PATIENT AGE 6 MONTHS TO < 6 YEARS WHO HAS ONE CFTR MUTATION RESPONSIVE TO IVACAFTOR BASED ON CLINICAL AND/OR IN VITRO ASSAY DATA USING A SOLID COMPOSITION COMPRISING AMORPHOUS (LESS THAN ABOUT 30% CRYSTALLINE) IVACAFTOR
- U-2531 TREATMENT OF CF IN A PATIENT AGE 6 MONTHS TO <6 YEARS WHO HAS ONE MUTATION IN THE CFTR GENE THAT IS RESPONSIVE TO IVACAFTOR BASED ON CLINICAL AND/OR IN VITRO ASSAY DATA USING THE COMPOSITION RECITED IN CLAIM 1 OF US 10272046
- U-2532 TREATMENT OF CHRONIC HEPATITIS C VIRUS (HCV) GENOTYPE 1, 2, 3, 4, OR 6 IN ADULT AND PEDIATRIC PATIENTS 12 YEARS AND OLDER OR WEIGHING AT LEAST 45 KG
- U-2533 A METHOD OF TREATING A CANCER CHARACTERIZED BY AN IDH1 MUTATION WHERE THE CANCER IS RELAPSED OR REFRACTORY ACUTE MYELOID LEUKEMIA (AML)
- U-2534 A METHOD OF TREATING A CANCER CHARACTERIZED BY AN IDH1 MUTATION WHERE THE CANCER IS NEWLY DIAGNOSED ACUTE MYELOID LEUKEMIA (AML)
- U-2535 USE IN COMBINATION WITH METHYLPREDNISOLONE FOR THE TREATMENT OF PATIENTS WITH PROSTATE CANCER
- U-2536 FOR TREATMENT OF STEROID-REFRACTORY ACUTE GRAFT-VERSUS-HOST DISEASE
- U-2537 TREATMENT OF ADULT PATIENTS WITH CHRONIC LYMPHOCYTIC LEUKEMIA (CLL) OR SMALL LYMPHOCYTIC LYMPHOMA (SLL)
- U-2538 TREATMENT OF ADULT PATIENTS WITH CHRONIC LYMPHOCYTIC LEUKEMIA (CLL) OR SMALL LYMPHOCYTIC LEUKEMIA (SLL) IN COMBINATION WITH A GA101 ANTIBODY SUCH AS OBINUTUZUMAB FOR ONE OR MORE DOSING PERIODS, WHEREIN THE CLL OR SLL IS A CD20-EXPRESSING CANCER
- U-2539 IN COMBINATION WITH FULVESTRANT FOR TREATMENT OF POSTMENOPAUSAL WOMEN, AND MEN, WITH HR-POSITIVE, HER-2-NEGATIVE, PIK3CA-MUTATED, ADVANCED OR METASTATIC BREAST CANCER
- U-2540 TREATMENT OF HORMONE RECEPTOR POSITIVE ADVANCED BREAST CANCER IN POSTMENOPAUSAL WOMEN
- U-2541 REDUCING THE RATE OF CARDIOVASCULAR DEATH, MYOCARDIAL INFARCTION (MI), AND STROKE IN A PATIENT RECEIVING 75-100 MG ASPIRIN DAILY WITH A HISTORY OF MI BY ADMINISTERING 60 MG TICAGRELOR TWICE DAILY
- U-2542 REDUCING THE RATE OF CARDIOVASCULAR DEATH, MYOCARDIAL INFARCTION, AND STROKE IN A PATIENT RECEIVING 75-100 MG ASPIRIN DAILY AND HAVING OR WHO HAD ACUTE CORONARY SYNDROME BY ADMINISTERING 60 MG TICAGRELOR TWICE DAILY
- U-2543 TREATMENT OF SCHIZOPHRENIA WITH CARIPRAZINE
- U-2544 TREATMENT OF ACUTE MANIC OR MIXED EPISODES ASSOCIATED WITH BIPOLAR I DISORDER WITH CARIPRAZINE
- U-2545 TREATMENT OF DEPRESSIVE EPISODES ASSOCIATED WITH BIPOLAR I DISORDER (BIPOLAR DEPRESSION) WITH CARIPRAZINE
- U-2546 USE FOR THE MAINTENANCE TREATMENT OF PATIENTS WITH CHRONIC OBSTRUCTIVE PULMONARY DISEASE (COPD), INCLUDING CHRONIC BRONCHITIS AND/OR EMPHYSEMA
- U-2547 METHOD OF PROVIDING CONTRACEPTION IN A WOMAN HAVING A BMI OF 25 KG/M2 OR MORE WITH RESULTANT LIMITED BLEEDING EVENTS PER TREATMENT CYCLE
- U-2548 TO IMPROVE WAKEFULNESS IN ADULT PATIENTS WITH EXCESSIVE DAYTIME SLEEPINESS ASSOCIATED WITH NARCOLEPSY OR OBSTRUCTIVE SLEEP APNEA (OSA)
- U-2549 CONTROL OF SERUM PHOSPHORUS LEVELS
- U-2550 USE OF REVLIMID (LENALIDOMIDE) FOR THE TREATMENT OF PREVIOUSLY TREATED FOLLICULAR LYMPHOMA IN COMBINATION WITH A RITUXIMAB PRODUCT
- U-2551 USE OF REVLIMID (LENALIDOMIDE) FOR THE TREATMENT OF PREVIOUSLY TREATED MARGINAL ZONE LYMPHOMA IN COMBINATION WITH A RITUXIMAB PRODUCT
- U-2552 METHOD OF TREATING POSTPARTUM DEPRESSION
- U-2553 PREVENTION OF PREGNANCY IN FEMALES OF REPRODUCTIVE AGE
- U-2554 TREATMENT OF MILD TO MODERATE ACTIVE CROHN'S DISEASE INVOLVING THE ILEUM AND/OR THE ASCENDING COLON

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

- U-2555 A METHOD OF TREATING IRON DEFICIENCY ANEMIA IN ADULT PATIENTS WHO HAVE INTOLERANCE TO ORAL IRON OR HAVE HAD UNSATISFACTORY RESPONSE TO ORAL IRON BY INTRAVENOUSLY ADMINISTERING FERRIC CARBOXYMALTOSATE TO PROVIDE AT LEAST ABOUT 0.6 G OF ELEMENTAL IRON
- U-2556 METHOD OF TREATING IRON DEFICIENCY ANEMIA IN ADULTS WHO HAVE INTOLERANCE TO OR HAVE HAD UNSATISFACTORY RESPONSE TO ORAL IRON ASSOCIATED WITH HEAVY UTERINE BLEEDING OR A GASTROINTESTINAL DISORDER BY INTRAVENOUSLY ADMINISTERING FERRIC CARBOXYMALTOSATE
- U-2557 A METHOD OF TREATING IRON DEFICIENCY ANEMIA IN ADULT PATIENTS WHO HAVE NON-DIALYSIS DEPENDENT CHRONIC KIDNEY DISEASE BY INTRAVENOUSLY ADMINISTERING FERRIC CARBOXYMALTOSATE TO PROVIDE AT LEAST ABOUT 0.6 GRAMS OF ELEMENTAL IRON
- U-2558 TREATMENT OF PATIENTS WITH LOCALLY ADVANCED METASTATIC BREAST CANCER AFTER FAILURE OF PRIOR CHEMOTHERAPY
- U-2559 USE IN COMBINATION WITH DOXORUBICIN AND CYCLOPHOSPHAMIDE FOR ADJUVANT TREATMENT OF PATIENTS WITH OPERABLE NODE-POSITIVE BREAST CANCER
- U-2560 TREATMENT OF PATIENTS WITH LOCALLY ADVANCED OR METASTATIC NON-SMALL CELL LUNG CANCER AFTER FAILURE OF PRIOR PLATINUM-BASED CHEMOTHERAPY
- U-2561 USE IN COMBINATION WITH CISPLATIN FOR TREATMENT OF UNRESECTABLE, LOCALLY ADVANCED OR METASTATIC NON-SMALL CELL LUNG CANCER WITHOUT PRIOR CHEMOTHERAPY TREATMENT
- U-2562 TREATMENT OF PATIENTS WITH ANDROGEN INDEPENDENT (HORMONE REFRACTORY) METASTATIC PROSTATE CANCER IN COMBINATION WITH PREDNISONE
- U-2563 TREATMENT OF ADVANCED GASTRIC ADENOCARCINOMA IN COMBINATION WITH CISPLATIN AND FLUOROURACIL IN PATIENTS THAT HAVE NOT RECEIVED PRIOR CHEMOTHERAPY
- U-2564 TREATMENT OF PATIENTS WITH LOCALLY ADVANCED SQUAMOUS CELL CARCINOMA OF THE HEAD AND NECK IN COMBINATION WITH CISPLATIN AND FLUOROURACIL
- U-2565 TREATMENT OF HOSPITAL-ACQUIRED BACTERIAL PNEUMONIA
- U-2566 TREATMENT OF VENTILATOR-ASSOCIATED BACTERIAL PNEUMONIA
- U-2567 ONCE DAILY TOPICAL TREATMENT OF PERSISTENT FACIAL ERYTHEMA ASSOCIATED WITH ROSACEA IN ADULTS WITH 1% OXYMETAZOLINE HYDROCHLORIDE CREAM, WHERE THE PATIENT EXPERIENCES NO REBOUND OR WORSENING OF FACIAL ERYTHEMA POST-TREATMENT
- U-2568 TREATMENT OF HYPOACTIVE SEXUAL DESIRE DISORDER (HSDD)
- U-2569 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 6 YEARS AND OLDER, WHO ARE HOMOZYGOUS FOR THE F508DEL MUTATION OR HAVE AT LEAST ONE CFTR MUTATION THAT IS RESPONSIVE TO TEZACAFTOR/IVACAFTOR, WITH AN EFFECTIVE AMOUNT OF TEZACAFTOR AND IVACAFTOR
- U-2570 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGES 6 AND OLDER, WHO HAVE TWO COPIES OF THE F508DEL MUTATION OR ONE F508DEL MUTATION AND A CFTR MUTATION PREDICTED TO BE RESPONSIVE TO TEZACAFTOR/IVACAFTOR, WITH THE COMPOSITION OF CLAIM 1 OF US 10,206,877
- U-2571 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGES 6 AND OLDER, WHO HAVE TWO COPIES OF THE F508DEL MUTATION OR ONE F508DEL MUTATION AND A CFTR MUTATION PREDICTED TO BE RESPONSIVE TO TEZACAFTOR/IVACAFTOR, WITH THE COMPOSITION OF CLAIM 1 OF US 10,081,621
- U-2572 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS 6 YEARS AND OLDER, WITH A F508DEL OR G551D CFTR GENE MUTATION AND A A455E, 2789+5G->, OR 3849+10KBC->T MUTATION, COMPRISING CONCURRENT COADMINISTRATION OF THE COMPOSITIONS OF CLAIM 1 OF U.S. PATENT 10058546
- U-2573 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 6 YEARS AND OLDER, WHO ARE HOMOZYGOUS FOR THE F508DEL MUTATION OR HETEROZYGOUS FOR F508DEL AND A SECOND CFTR MUTATION PREDICTED TO BE RESPONSIVE TO TEZACAFTOR/IVACAFTOR, WITH TEZACAFTOR AND IVACAFTOR
- U-2574 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 6 AND OLDER, WHO ARE HOMOZYGOUS FOR THE F508DEL MUTATION OR HAVE AT LEAST ONE CFTR GENE MUTATION THAT IS RESPONSIVE TO TEZACAFTOR/IVACAFTOR, WITH TEZACAFTOR AND IVACAFTOR
- U-2575 TREATING CYSTIC FIBROSIS PATIENTS AGES 6 AND OLDER, WHO ARE HOMOZYGOUS FOR F508DEL OR HAVE AT LEAST 1 CFTR GENE MUTATION RESPONSIVE TO TEZACAFTOR/IVACAFTOR, WITH TEZACAFTOR AND A SOLID COMPOSITION COMPRISING AMORPHOUS (<30% CRYSTALLINE) IVACAFTOR
- U-2576 TREATMENT OF COMMUNITY ACQUIRED BACTERIAL PNEUMONIA

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

- U-2577 TREATMENT OF THROMBOCYTOPENIA IN AN ADULT PATIENT WITH CHRONIC IMMUNE THROMBOCYTOPENIA WHO HAS HAD AN INSUFFICIENT RESPONSE TO A PREVIOUS TREATMENT
- U-2578 TREATMENT OF THROMBOCYTOPENIA IN AN ADULT PATIENT WITH CHRONIC LIVER DISEASE WHO IS SCHEDULED TO UNDERGO A PROCEDURE
- U-2579 REDUCTION IN A SUBJECT'S RISK OF EXPERIENCING A BREAKTHROUGH OVERT HEPATIC ENCEPHALOPATHY (HE) EPISODE
- U-2580 A METHOD OF TREATING TYPE 2 DIABETES COMPRISING ADMINISTERING SEMAGLUTIDE ONCE WEEKLY IN A AMOUNT OF 1.0 MG TO A SUBJECT IN NEED THEREOF
- U-2581 TREATING HYPOTENSION WITH ABOUT 20 NG/KG/MIN TO ABOUT 40 NG/KG/MIN ANGIOTENSIN II IN A HUMAN SUBJECT HAVING SEPTIC SHOCK
- U-2582 FOR THE ORAL PREVENTION/PROPHYLAXIS OF MALARIA IN ADULTS, COMPRISING A THREE-PHASE DOSING REGIMEN CONSISTING OF A LOADING/INITIAL DOSE, A MAINTENANCE/EXPOSURE DOSE, AND A TERMINAL/POST-EXPOSURE DOSE
- U-2583 TREATMENT OF BACTERIAL VAGINOSIS IN ADULT WOMEN
- U-2584 XPROVIO IS INDICATED IN COMBINATION WITH DEXAMETHASONE TO TREAT RELAPSED OR REFRACTORY MULTIPLE MYELOMA (REFRACTORY TO AT LEAST AN ANTI-CD38 MAB, 2 PROTEASOME INHIBITORS AND 2 IMMUNOMODULATORY AGENTS) IN ADULTS WHO RECEIVED AT LEAST 4 PRIOR THERAPIES
- U-2585 TREATMENT OF PARENTERAL NUTRITION-ASSOCIATED CHOLESTASIS IN PATIENTS UNDER THE AGE OF 12
- U-2586 TREATMENT OF COMPLICATED URINARY TRACT INFECTIONS, INCLUDING PYELONEPHRITIS (CUTI)
- U-2587 TREATMENT OF COMPLICATED INTRA-ABDOMINAL INFECTIONS (CIAI)
- U-2588 AS AN ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS IN COMBINATION WITH DAPAGLIFLOZIN AND METFORMIN
- U-2589 AS AN ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS IN COMBINATION WITH BASAL INSULIN OR BASAL INSULIN PLUS METFORMIN
- U-2590 AS AN ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS IN COMBINATION WITH METFORMIN, A SULFONYLUREA, A THIAZOLIDINEDIONE, OR COMBINATION OF ANY TWO OF THESE THERAPIES
- U-2591 LOWERING PLASMA GLUCAGON IN ADULTS WITH TYPE 2 DIABETES MELLITUS BY ADMINISTERING EXENATIDE AS AN ADJUNCT TO DIET AND EXERCISE
- U-2592 IMPROVING GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS BY ADMINISTERING A SUSTAINED-RELEASE EXENATIDE FORMULATION AS AN ADJUNCT TO DIET AND EXERCISE
- U-2593 IMPROVING GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS BY ADMINISTERING AN EXENATIDE FORMULATION ONCE WEEKLY AS AN ADJUNCT TO DIET AND EXERCISE TO ACHIEVE A MEAN STEADY STATE PLASMA CONCENTRATION OF EXENATIDE AT LEAST 170 PG/ML
- U-2594 REDUCING FASTING PLASMA GLUCOSE IN ADULTS WITH TYPE 2 DIABETES MELLITUS BY ADMINISTERING AN EXENATIDE FORMULATION ONCE WEEKLY AS AN ADJUNCT TO DIET AND EXERCISE TO ACHIEVE A MEAN STEADY STATE PLASMA CONCENTRATION OF EXENATIDE AT LEAST 170 PG/ML
- U-2595 REDUCING BODY WEIGHT IN ADULTS WITH TYPE 2 DIABETES MELLITUS BY ADMINISTERING AN EXENATIDE FORMULATION ONCE WEEKLY AS AN ADJUNCT TO DIET AND EXERCISE TO ACHIEVE A MEAN STEADY STATE PLASMA CONCENTRATION OF EXENATIDE AT LEAST 170 PG/ML
- U-2596 REDUCING HBA1C IN ADULTS WITH TYPE 2 DIABETES MELLITUS BY ADMINISTERING AN EXENATIDE FORMULATION ONCE WEEKLY AS AN ADJUNCT TO DIET AND EXERCISE TO ACHIEVE A MEAN STEADY STATE PLASMA CONCENTRATION OF EXENATIDE AT LEAST 170 PG/ML
- U-2597 AS AN ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS IN COMBINATION WITH DAPAGLIFLOZIN AS ADD-ON TO METFORMIN
- U-2598 IMPROVING GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS BY ADMINISTERING AN INJECTABLE SUSTAINED RELEASE FORMULATION OF EXENATIDE AS AN ADJUNCT TO DIET AND EXERCISE
- U-2599 IMPROVING GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS BY ADMINISTERING AN EXENATIDE FORMULATION AS AN ADJUNCT TO DIET AND EXERCISE TO PROVIDE A RELEASE PROFILE HAVING A RATIO OF C-MAX TO C-AVG OF ABOUT 3 OR LESS

PATENT AND EXCLUSIVITY TERMS

ADB 162 of 225

PATENT USE

U-2600 IMPROVING GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS BY ADMINISTERING A PRE-MIXED EXENATIDE FORMULATION AS AN ADJUNCT TO DIET AND EXERCISE

U-2601 STIMULATING INSULIN RELEASE IN ADULTS WITH TYPE 2 DIABETES MELLITUS BY ADMINISTERING A PRE-MIXED EXENATIDE FORMULATION AS AN ADJUNCT TO DIET AND EXERCISE

U-2602 DELAYING GASTRIC EMPTYING IN ADULTS WITH TYPE 2 DIABETES MELLITUS BY ADMINISTERING A PRE-MIXED EXENATIDE FORMULATION AS AN ADJUNCT TO DIET AND EXERCISE

U-2603 METHOD OF TREATING IRON DEFICIENCY

U-2604 TREATMENT OF SEVERE HYPOGLYCEMIA IN PATIENTS WITH DIABETES

U-2605 TREATMENT OF PATIENTS WITH NON-METASTATIC CASTRATION RESISTANT PROSTATE CANCER

U-2606 TREATMENT OF ADULT PATIENTS WITH SYMPTOMATIC TENOSYNOVIAL GIANT CELL TUMOR (TGCT) ASSOCIATED WITH SEVERE MORBIDITY OR FUNCTIONAL LIMITATIONS AND NOT AMENABLE TO IMPROVEMENT WITH SURGERY

U-2607 TREATMENT OF ADULT PATIENTS WITH INTERMEDIATE-2 OR HIGH-RISK PRIMARY OR SECONDARY MYELOFIBROSIS

U-2608 METHOD OF TREATING SCHIZOPHRENIA

U-2609 A METHOD FOR INDUCING A REGIONAL ANAESTHESIA VIA INTRATHECAL ADMINISTRATION OF A PATENTED PRESERVATIVE FREE SOLUTION FOR INJECTION (WITH A SPECIFIC COMPOSITION, PH, OSMOLALITY AND DENSITY) CONTAINING 9-11 MG/ML CHLOROPROCAINE HCL

U-2610 TREATMENT OF COMPLICATED INTRA-ABDOMINAL INFECTION IN PATIENTS WITH END-STAGE RENAL DISEASE ON HEMODIALYSIS

U-2611 TREATMENT OF COMPLICATED URINARY TRACT INFECTION IN PATIENTS WITH END-STAGE RENAL DISEASE ON HEMODIALYSIS

U-2612 TREATMENT OF RELAPSING FORMS OF MULTIPLE SCLEROSIS (MS), TO INCLUDE CLINICALLY ISOLATED SYNDROME, RELAPSING-REMITTING DISEASE, AND ACTIVE SECONDARY PROGRESSIVE DISEASE, IN PATIENTS 10 YEARS OF AGE AND OLDER

U-2613 TREATMENT OF RELAPSING-REMITTING SCLEROSIS (MS)

U-2614 TREATMENT OF MODERATE TO SEVERE DYSpareunia

U-2615 TREATMENT OF NON-24 HOUR SLEEP-WAKE DISORDER BY AVOIDING THE ADMINISTRATION OF TASIMELTEON WITH FOOD

U-2616 TREATMENT OF ADULTS WITH MODERATELY TO SEVERELY ACTIVE RHEUMATOID ARTHRITIS WHO HAVE HAD AN INADEQUATE RESPONSE OR INTOLERANCE TO METHOTREXATE

U-2617 TREATMENT OF ROS1-POSITIVE NON-SMALL CELL LUNG CANCER

U-2618 TREATMENT OF SOLID TUMORS THAT HAVE A NEUROTROPHIC TYROSINE RECEPTOR KINASE (NTRK) GENE FUSION

U-2619 TREATMENT OF ADULTS WITH COMMUNITY-ACQUIRED BACTERIAL PNEUMONIA CAUSED BY SUSCEPTIBLE MICROORGANISMS

U-2620 USE OF NINTEDANIB FOR SLOWING THE RATE OF DECLINE IN PULMONARY FUNCTION IN PATIENTS WITH SYSTEMIC SCLEROSIS-ASSOCIATED INTERSTITIAL LUNG DISEASE (SSC-ILD)

U-2621 MODIFIED DOSING REGIMEN FOR THE MANAGEMENT OF MILD TO MODERATE PAIN

U-2622 TREATMENT OF ACUTE UNCOMPLICATED INFLUENZA IN PATIENTS 2 YEARS AND OLDER

U-2623 A METHOD OF REDUCING OFF TIME FROM L-DOPA THERAPY, COMPRISING ADMINISTERING, TO A HUMAN PATIENT WITH PARKINSON'S DISEASE, AN EFFECTIVE AMOUNT OF ISTRADEFYLLINE, WHEREIN THE PATIENT CURRENTLY RECEIVES SAID L-DOPA THERAPY

U-2624 TREATMENT OF METASTATIC CASTRATION-SENSITIVE PROSTATE CANCER (MCSPC)

U-2625 TOPICAL TREATMENT OF PLAQUE PSORIASIS IN ADULTS

U-2626 METHOD OF TREATING IRRITABLE BOWEL SYNDROME WITH CONSTIPATION BY ADMINISTERING TENAPANOR

U-2627 TOPICAL TREATMENT OF PLAQUE PSORIASIS IN PATIENTS 12 YEARS AND OLDER

U-2628 METHOD OF TREATING TYPE 2 DIABETES MELLITUS

U-2629 TREATMENT OF HIV-1 INFECTION IN ADULT PATIENTS AS A REPLACEMENT THERAPY IN VIROLOGICALLY SUPPRESSED ADULTS WITH NO HISTORY OF TREATMENT FAILURE AND NO KNOWN SUBSTITUTIONS ASSOCIATED WITH RESISTANCE TO THE INDIVIDUAL COMPONENTS OF DELSTRIGO

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

- U-2630 FOR USE IN COMBINATION WITH OTHER ANTIRETROVIRAL AGENTS FOR THE TREATMENT OF HIV-1 AS A REPLACEMENT THERAPY IN VIROLOGICALLY SUPPRESSED ADULTS WITH NO HISTORY OF TREATMENT FAILURE AND NO KNOWN SUBSTITUTIONS ASSOCIATED WITH RESISTANCE TO DORAVIRINE
- U-2631 TREATMENT OF COMPLICATED URINARY TRACT INFECTION
- U-2632 REDUCTION OF RISK OF END STAGE KIDNEY DISEASE, DOUBLING OF SERUM CREATININE, CARDIOVASCULAR DEATH, AND HOSPITALIZATION FOR HEART FAILURE IN THE TREATMENT OF TYPE 2 DIABETES MELLITUS PATIENTS
- U-2633 TREATMENT OF ANAPLASTIC LYMPHOMA KINASE (ALK)-POSITIVE METASTATIC NON-SMALL CELL LUNG CANCER, PROGRESSED ON: CRIZOTINIB + AT LEAST 1 OTHER ALK INHIBITOR FOR METASTATIC DISEASE; OR ALECTINIB, OR CERITINIB AS FIRST ALK INHIBITOR FOR METASTATIC DISEASE.
- U-2634 METHOD OF TREATMENT IN PATIENTS WITH CONCOMITANT ANGIOEDEMA
- U-2635 TREATMENT OF ACUTE URTICARIA
- U-2636 METHOD OF INCREASING PEAK PLASMA OR ONSET OF PLASMA CONCENTRATION BY INTRAVENOUS INJECTION IN INDIVIDUALS IN NEED OF TREATMENT FOR ACUTE URTICARIA
- U-2637 TREATMENT OF PATIENTS WITH AN OVERACTIVE BLADDER WITH SYMPTOMS OF URINARY FREQUENCY, URGENCY, OR URGE INCONTINENCE WITH A SINGLE UNIT DOSE OF 10% OXYBUTYNIN CHLORIDE GEL
- U-2638 INCREASE PAIN-FREE LIGHT EXPOSURE IN ADULT PATIENTS WITH A HISTORY OF PHOTOTOXIC REACTIONS FROM ERYTHROPOIETIC PROTOPORPHYRIA (EPP)
- U-2639 METHOD OF ACTIVATING RARGAMMA RECEPTOR
- U-2640 PROPHYLAXIS OF VENOUS THROMBOEMBOLISM IN ACUTELY ILL MEDICAL PATIENTS AT RISK FOR THROMBOEMBOLIC COMPLICATIONS NOT AT HIGH RISK OF BLEEDING
- U-2641 PROPHYLAXIS OF VENOUS THROMBOEMBOLISM IN ACUTELY ILL MEDICAL PATIENTS AT RISK FOR THROMBOEMBOLIC COMPLICATIONS NOT AT HIGH RISK OF BLEEDING WITH ONCE DAILY, RAPID-RELEASE TABLET ADMINISTERED FOR AT LEAST 5 CONSECUTIVE DAYS
- U-2642 METHOD OF TREATING CANCER BY DETECTING A CREATININE CLEARANCE OF A PATIENT AND ADMINISTERING LONSURF
- U-2643 TREATMENT OF IRRITABLE BOWEL SYNDROME WITH DIARRHEA (IBS-D) IN ADULTS 65 YEARS OF AGE OR OLDER AND SYMPTOMS THEREOF
- U-2644 TREATMENT OF IRRITABLE BOWEL SYNDROME WITH DIARRHEA (IBS-D) IN ADULTS 65 YEARS OF AGE OR OLDER
- U-2645 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 12 YEARS AND OLDER WHO HAVE AT LEAST ONE F508DEL MUTATION IN THE CFTR GENE WITH ELEXACAFTOR, TEZACAFTOR, AND IVACAFTOR
- U-2646 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 12 YEARS AND OLDER WHO HAVE ONE F508DEL MUTATION AND ONE R117H MUTATION IN THE CFTR GENE WITH ELEXACAFTOR, TEZACAFTOR, AND IVACAFTOR
- U-2647 TREATMENT OF NON-NODULAR ACNE VULGARIS
- U-2648 TREATMENT OF A MODERATE TO MILD CLINICAL PHENOTYPE OF CF IN PATIENTS AGED 12 YEARS AND OLDER WHO HAVE AT LEAST ONE F508DEL MUTATION IN THE CFTR GENE WITH ELEXACAFTOR, TEZACAFTOR, AND IVACAFTOR
- U-2649 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 12 YEARS AND OLDER WHO HAVE AT LEAST ONE F508DEL MUTATION IN THE CFTR GENE WITH A COMPOSITION COMPRISING ELEXACAFTOR, TEZACAFTOR, AND IVACAFTOR; AND ANOTHER COMPOSITION COMPRISING IVACAFTOR
- U-2650 TREATMENT OF CF IN PATIENTS AGED 12 YEARS AND OLDER WHO HAVE AT LEAST ONE F508DEL MUTATION IN THE CFTR GENE USING A SOLID COMPOSITION COMPRISING ELEXACAFTOR, TEZACAFTOR, AMORPHOUS IVACAFTOR, AND LESS THAN ABOUT 30% CRYSTALLINE IVACAFTOR
- U-2651 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 12 YEARS AND OLDER WHO HAVE AT LEAST ONE F508DEL MUTATION IN THE CFTR GENE WITH AN EFFECTIVE AMOUNT OF A PHARMACEUTICAL COMPOSITION COMPRISING ELEXACAFTOR, TEZACAFTOR, AND IVACAFTOR
- U-2652 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 12 YEARS AND OLDER WHO HAVE AT LEAST ONE F508DEL MUTATION IN THE CFTR GENE WITH A COMPOSITION ACCORDING TO CLAIM 1 OF US 10081621
- U-2653 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 12 YEARS AND OLDER WHO HAVE AT LEAST ONE F508DEL MUTATION IN THE CFTR GENE WITH AN EFFECTIVE AMOUNT OF

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

ELEXACAFTOR, TEZACAFTOR, AND IVACAFTOR

- U-2654 TREATMENT OF REFRACTORY CHRONIC GRAFT-VERSUS-HOST DISEASE
- U-2655 A METHOD OF TREATMENT OF ADVANCED OVARIAN, FALLOPIAN TUBE, OR PRIMARY PERITONEAL CANCER ASSOCIATED WITH HOMOLOGOUS RECOMBINATION DEFICIENCY (HRD) POSITIVE STATUS
- U-2656 TREATMENT OF ADULT PATIENTS WITH ACTIVE PSORIATIC ARTHRITIS
- U-2657 TREATMENT OF PATIENTS WITH MODERATE TO SEVERE PLAQUE PSORIASIS WHO ARE CANDIDATES FOR PHOTOTHERAPY OR SYSTEMIC THERAPY
- U-2658 TREATMENT OF ADULT PATIENTS WITH ORAL ULCERS ASSOCIATED WITH BEHCET'S DISEASE
- U-2659 TREATMENT OF ADULT PATIENTS WITH ORAL ULCERS ASSOCIATED WITH BEHCET'S DISEASE USING A DOSAGE TITRATION SCHEDULE
- U-2660 TREATMENT OF H. PYLORI INFECTION IN ADULTS
- U-2661 CHRONIC WEIGHT MANAGEMENT IN ADULT PATIENTS USING AN EXTENDED RELEASE TABLET CONTAINING LORCARSERIN HYDROCHLORIDE HEMIHYDRATE
- U-2662 USE OF CALCIPOTRIENE FOAM FOR THE TOPICAL TREATMENT OF PLAQUE PSORIASIS IN PATIENTS AGED 4 YEARS AND OLDER
- U-2663 USE IN SONOHYSTEROSALPINOGRAPHY TO ASSESS FALLOPIAN TUBE PATENCY
- U-2664 TREATMENT OF DUCHENNE MUSCULAR DYSTROPHY (DMD) IN PATIENTS HAVING A MUTATION OF THE DMD GENE THAT IS AMENABLE TO EXON 51 SKIPPING BY INDUCING SKIPPING OF EXON 51
- U-2665 TREATMENT OF CHRONIC GRAFT VERSUS HOST DISEASE AFTER FAILURE OF ONE OR MORE LINES OF SYSTEMIC THERAPY
- U-2666 TREATMENT OF ADULT PATIENTS WITH CHRONIC LYMPHOCYTIC LEUKEMIA
- U-2667 TREATMENT OF ADULT PATIENTS WITH SMALL LYMPHOCYTIC LEUKEMIA
- U-2668 TREATMENT OF ADULT PATIENTS WITH PREVIOUSLY UNTREATED CHRONIC LYMPHOCYTIC LEUKEMIA IN COMBINATION WITH OBINUTUZUMAB
- U-2669 TREATMENT OF ADULT PATIENTS WITH PREVIOUSLY UNTREATED SMALL LYMPHOCYTIC LEUKEMIA IN COMBINATION WITH OBINUTUZUMAB
- U-2670 TREATMENT OF ADULT PATIENTS WITH CHRONIC LYMPHOCYTIC LEUKEMIA OR SMALL LYMPHOCYTIC LEUKEMIA
- U-2671 TREATMENT OF ADULT PATIENTS WITH PREVIOUSLY UNTREATED CHRONIC LYMPHOCYTIC LEUKEMIA OR SMALL LYMPHOCYTIC LEUKEMIA IN COMBINATION WITH OBINUTUZUMAB
- U-2672 TREATMENT OF ACUTE HEPATIC PORPHYRIA
- U-2673 TREATMENT OF DUCHENNE MUSCULAR DYSTROPHY (DMD) IN PATIENTS HAVING A MUTATION OF THE DMD GENE THAT IS AMENABLE TO EXON 51 SKIPPING BY CORRECTING A DEFECTIVE GENE FOR DYSTROPHIN
- U-2674 TREATMENT OF DUCHENNE MUSCULAR DYSTROPHY (DMD) IN PATIENTS HAVING A MUTATION OF THE DMD GENE THAT IS AMENABLE TO EXON 51 SKIPPING BY RESTORING OR INCREASING FUNCTIONAL DYSTROPHIN PROTEIN PRODUCTION
- U-2675 TREATMENT OF DUCHENNE MUSCULAR DYSTROPHY (DMD) IN PATIENTS HAVING A CONFIRMED MUTATION OF THE DMD GENE THAT IS AMENABLE TO EXON 53 SKIPPING
- U-2676 TREATMENT OF SICKLE CELL DISEASE BY ADMINISTERING VOXELOTOR, AS RECITED IN CLAIM 1
- U-2677 PROPHYLAXIS OF ORGAN REJECTION IN PATIENTS CONVERTED FROM TACROLIMUS IMMEDIATE-RELEASE FORMULATIONS
- U-2678 PROPHYLAXIS OF ORGAN REJECTION IN DE NOVO TRANSPLANT PATIENT
- U-2679 TREATING LOW BLOOD PRESSURE WITH ANGIOTENSIN II AT AN INITIAL RATE OF ABOUT 20 NG/KG/MIN AND TITRATING DOWN TO ACHIEVE AND/OR MAINTAIN A MAP OF ABOUT 65 MM HG OR ABOVE
- U-2680 TREATING LOW BLOOD PRESSURE WITH ANGIOTENSIN II WITH AN INITIAL RATE OF ABOUT 5 NG/KG/MIN TO ABOUT 20 NG/KG/MIN IN A SUBJECT HAVING REFRACTORY HYPOTENSION OR SEVERE HYPOTENSION
- U-2681 TREATING LOW BLOOD PRESSURE WITH ANGIOTENSIN II WITH AN INITIAL RATE OF ABOUT 5 NG/KG/MIN TO ABOUT 20 NG/KG/MIN IN A SUBJECT HAVING REFRACTORY HYPOTENSION OR SEVERE HYPOTENSION, AND TITRATING THE RATE UP
- U-2682 TREATMENT OF ADULT PATIENTS WITH CHRONIC LYMPHOCYTIC LEUKEMIA BY ADMINISTERING 100MG OF ACALABRUTINIB TWICE DAILY

PATENT AND EXCLUSIVITY TERMS

ADB 165 of 225

PATENT USE

- U-2683 TREATMENT OF ADULT PATIENTS WITH SMALL LYMPHOCYTIC LEUKEMIA BY ADMINISTERING 100MG OF ACALABRUTINIB TWICE DAILY
- U-2684 TREATMENT OF ADULT PATIENTS WITH PREVIOUSLY UNTREATED CHRONIC LYMPHOCYTIC LEUKEMIA BY ADMINISTERING 100 MG OF ACALABRUTINIB TWICE DAILY IN COMBINATION WITH OBINUTUZUMAB
- U-2685 TREATMENT OF ADULT PATIENTS WITH PREVIOUSLY UNTREATED SMALL LYMPHOCYTIC LEUKEMIA BY ADMINISTERING 100 MG OF ACALABRUTINIB TWICE DAILY IN COMBINATION WITH OBINUTUZUMAB
- U-2686 TREATMENT OF ADULT PATIENTS WITH CHRONIC LYMPHOCYTIC LEUKEMIA OR SMALL LYMPHOCYTIC LEUKEMIA BY ADMINISTERING 100 MG OF ACALABRUTINIB TWICE DAILY
- U-2687 TREATMENT OF ADULT PATIENTS WITH PREVIOUSLY UNTREATED CHRONIC LYMPHOCYTIC LEUKEMIA OR SMALL LYMPHOCYTIC LEUKEMIA IN COMBINATION WITH OBINUTUZUMAB BY ADMINISTERING 100 MG OF ACALABRUTINIB TWICE DAILY
- U-2688 USE OF VASCEPA TO LOWER TRIGLYCERIDES AND LDL-C IN AN ADULT PATIENT WITH ELEVATED TRIGLYCERIDE (TG) LEVELS (ABOUT 200 MG/DL TO LESS THAN ABOUT 500 MG/DL) AND ON STATIN THERAPY
- U-2689 USE OF VASCEPA TO TREAT MIXED DYSLIPIDEMIA IN AN ADULT PATIENT WITH ELEVATED TRIGLYCERIDE (TG) LEVELS (\geq 150 MG/DL) AND ON STATIN THERAPY
- U-2690 USE OF VASCEPA TO LOWER TRIGLYCERIDES IN AN ADULT PATIENT WITH ELEVATED TRIGLYCERIDE (TG) LEVELS (ABOUT 200 MG/DL TO LESS THAN ABOUT 500 MG/DL) AND ON STATIN THERAPY
- U-2691 USE OF VASCEPA TO TREAT HYPERTRIGLYCERIDEMIA IN AN ADULT PATIENT WITH ELEVATED TRIGLYCERIDE (TG) LEVELS (\geq 150 MG/DL) AND ON STATIN THERAPY
- U-2692 USE OF VASCEPA TO REDUCE TRIGLYCERIDES IN AN ADULT PATIENT WITH ELEVATED TRIGLYCERIDE (TG) LEVELS (\geq 150 MG/DL) AND ON STATIN THERAPY
- U-2693 USE OF VASCEPA TO REDUCE TRIGLYCERIDES IN A MIXED DYSLIPIDEMIA ADULT PATIENT WITH ELEVATED TRIGLYCERIDE (TG) LEVELS (\geq 150 MG/DL) AND ON STATIN THERAPY
- U-2694 USE OF VASCEPA TO LOWER TRIGLYCERIDES IN A MIXED DYSLIPIDEMIA ADULT PATIENT WITH ELEVATED TRIGLYCERIDE (TG) LEVELS (ABOUT 200 MG/DL TO LESS THAN ABOUT 500 MG/DL) AND ON STATIN THERAPY
- U-2695 USE OF VASCEPA TO TREAT MIXED HYPERTRIGLYCERIDEMIA IN AN ADULT PATIENT WITH ELEVATED TRIGLYCERIDE (TG) LEVELS (\geq 150 MG/DL) AND ON STATIN THERAPY
- U-2696 USE OF VASCEPA AS AN ADJUNCT TO STATIN THERAPY TO REDUCE THE RISK OF CARDIOVASCULAR DEATH, CORONARY REVASCULARIZATION, AND UNSTABLE ANGINA IN AN ADULT PATIENT WITH ELEVATED TRIGLYCERIDE LEVELS (TG \geq 150 MG/DL TO ABOUT 500 MG/DL)
- U-2697 USE OF VASCEPA AS AN ADJUNCT TO STATIN THERAPY TO REDUCE THE RISK OF CARDIOVASCULAR DEATH AND/OR UNSTABLE ANGINA IN AN ADULT PATIENT WITH ELEVATED TRIGLYCERIDE LEVELS (TG \geq 150 MG/DL TO ABOUT 500 MG/DL)
- U-2698 USE OF VASCEPA AS AN ADJUNCT TO STATIN THERAPY TO REDUCE THE RISK OF CARDIOVASCULAR DEATH AND/OR CORONARY REVASCULARIZATION IN AN ADULT PATIENT WITH ELEVATED TRIGLYCERIDE LEVELS (TG \geq 150 MG/DL TO ABOUT 500 MG/DL)
- U-2699 USE OF VASCEPA AS AN ADJUNCT TO STATIN THERAPY TO REDUCE THE RISK OF A CARDIOVASCULAR EVENT (CORONARY REVASCULARIZATION, UNSTABLE ANGINA, STROKE AND/OR MYOCARDIAL INFARCTION) IN AN ADULT PATIENT WITH ELEVATED TRIGLYCERIDE LEVELS
- U-2700 USE OF VASCEPA TO REDUCE TRIGLYCERIDES IN AN ADULT PATIENT WITH ELEVATED TRIGLYCERIDE (TG) LEVELS (ABOUT 200 MG/DL TO LESS THAN ABOUT 500 MG/DL) AND ON ROSUVASTATIN THERAPY
- U-2701 USE OF VASCEPA AS AN ADJUNCT TO STATIN THERAPY TO REDUCE THE RISK OF CORONARY REVASCULARIZATION AND/OR UNSTABLE ANGINA IN AN ADULT PATIENT WITH ELEVATED TRIGLYCERIDE LEVELS (TG \geq 150 MG/DL TO ABOUT 500 MG/DL)
- U-2702 USE OF VASCEPA AS AN ADJUNCT TO STATIN THERAPY TO REDUCE THE RISK OF A CARDIOVASCULAR EVENT (CARDIOVASCULAR DEATH, CORONARY REVASCULARIZATION AND/OR UNSTABLE ANGINA) IN AN ADULT PATIENT WITH ELEVATED TRIGLYCERIDE LEVELS
- U-2703 USE OF VASCEPA AS AN ADJUNCT TO STATIN THERAPY TO REDUCE THE RISK OF A CV EVENT (CV DEATH, CORONARY REVASCULARIZATION, UNSTABLE ANGINA, STROKE AND/OR MYOCARDIAL INFARCTION) IN AN ADULT PATIENT WITH ELEVATED TRIGLYCERIDE LEVELS AND DIABETES MELLITUS
- U-2704 USE OF VASCEPA AS AN ADJUNCT TO STATIN THERAPY TO REDUCE THE RISK OF A CARDIOVASCULAR EVENT IN AN ADULT PATIENT WITH ELEVATED TRIGLYCERIDE LEVELS AND

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

AT LEAST ONE RISK FACTOR FOR CARDIOVASCULAR DISEASE

- U-2705 METHOD OF USING CAPSAICIN IN COMBINATION WITH A GEL COMPOSITION FOR REMOVAL OF CAPSAICIN FROM A TREATMENT AREA OR UNINTENDED AREA
- U-2706 USE OF VASCEPA AS AN ADJUNCT TO STATIN THERAPY TO REDUCE THE RISK OF ONSET AND/OR RECURRENCE OF CARDIOVASCULAR EVENTS IN A PATIENT WHO HAS ESCAPED THE UNSTABLE PERIOD AFTER CARDIOVASCULAR ANGIOPLASTY
- U-2707 USE OF VASCEPA AS AN ADJUNCT TO STATIN THERAPY TO REDUCE THE OCCURRENCE OF A CARDIOVASCULAR EVENT IN AN ADULT PATIENT WITH HYPERCHOLESTEROLEMIA
- U-2708 THE TREATMENT OF PATIENTS WITH METASTATIC CASTRATION-SENSITIVE PROSTATE CANCER
- U-2709 A METHOD OF TREATING IRON DEFICIENCY ANEMIA IN ADULT PATIENTS WHO HAVE INTOLERANCE TO ORAL IRON OR HAVE HAD UNSATISFACTORY RESPONSE TO ORAL IRON BY INTRAVENOUSLY ADMINISTERING FERRIC CARBOXYMALTOSE
- U-2710 A METHOD OF TREATING IRON DEFICIENCY ANEMIA IN ADULT PATIENTS WHO HAVE NON-DIALYSIS DEPENDENT CHRONIC KIDNEY DISEASE BY INTRAVENOUSLY ADMINISTERING FERRIC CARBOXYMALTOSE
- U-2711 A METHOD OF TREATING IRON DEFICIENCY ANEMIA IN ADULT PATIENTS WHO HAVE INTOLERANCE TO ORAL IRON OR HAVE HAD UNSATISFACTORY RESPONSE TO ORAL IRON BY INTRAVENOUSLY ADMINISTERING FERRIC CARBOXYMALTOSE TO PROVIDE 750 MG OF ELEMENTAL IRON
- U-2712 A METHOD OF TREATING IRON DEFICIENCY ANEMIA IN ADULT PATIENTS WHO HAVE NON-DIALYSIS DEPENDENT CHRONIC KIDNEY DISEASE BY INTRAVENOUSLY ADMINISTERING FERRIC CARBOXYMALTOSE TO PROVIDE 750 MG OF ELEMENTAL IRON
- U-2713 MODULATION OF 5-HYDROXYTRYPTAMINE 2 RECEPTOR ACTIVITY IN SCHIZOPHRENIA
- U-2714 TREATMENT OF SCHIZOPHRENIA WITH IMPROVEMENT IN RESIDUAL SYMPTOMS OF SCHIZOPHRENIA
- U-2715 TREATMENT OF SICKLE CELL DISEASE BY ADMINISTERING VOXELOTOR, AS RECITED IN CLAIM 2
- U-2716 MAINTENANCE TREATMENT OF DELETERIOUS OR SUSPECTED DELETERIOUS GBRCA-MUTATED METASTATIC PANCREATIC ADENOCARCINOMA WHOSE DISEASE HAS NOT PROGRESSED ON AT LEAST 16 WEEKS OF A FIRST-LINE PLATINUM-BASED CHEMOTHERAPY REGIMEN
- U-2717 ACUTE TREATMENT OF MIGRAINE WITH HEADACHE, WITH OR WITHOUT AURA IN ADULTS
- U-2718 ACUTE TREATMENT OF MIGRAINE WITH OR WITHOUT AURA IN ADULTS
- U-2719 TREATMENT OF RELAPSING REMITTING MULTIPLE SCLEROSIS BY DETERMINING VARICELLA ZOSTER VIRUS (VZV) STATUS AND VACCINATING PRIOR TO COMMENCING TREATMENT
- U-2720 ANTIMYCOTIC USES, SPECIFICALLY TREATMENT OF ONYCHOMYCOSIS; TOPICAL TREATMENT OF THE TOENAIL(S) DUE TO TRICHOPHYTON RUBRUM AND TRICHOPHYTON MENTAGROPHYTES
- U-2721 TOPICAL TREATMENT OF TINEA UNGUIUM BY USING AN APPLICATOR FOR APPLYING A SOLUTION FOR TREATING TINEA UNGUIUM TO AN AFFECTED PART OF A PATIENT
- U-2722 METHOD OF INTRAVENOUSLY ADMINISTERING A DILUTED CYSTEINE HYDROCHLORIDE SOLUTION TO A NEONATE IN NEED THEREOF
- U-2723 MAINTENANCE MONOTHERAPY TREATMENT OF BIPOLAR 1 DISORDER
- U-2724 A METHOD OF ORAL DELIVERY OF TREPROSTINIL COMPRISING ADMINISTERING AN ORAL OSMOTIC PHARMACEUTICAL DOSAGE FORM
- U-2725 A METHOD OF TREATING PULMONARY HYPERTENSION AND PULMONARY ARTERIAL HYPERTENSION BY ADMINISTERING AN ORAL OSMOTIC PHARMACEUTICAL DOSAGE FORM
- U-2726 TREATMENT OF UNRESECTABLE OR METASTATIC GASTROINTESTINAL STROMAL TUMOR (GIST) HARBORING A PLATELET-DERIVED GROWTH FACTOR RECEPTOR ALPHA (PDGFRA) EXON 18 MUTATION
- U-2727 NASAL ADMINISTRATION OF DIAZEPAM FOR TREATMENT OF INTERMITTENT, STEREOTYPIC EPISODES OF FREQUENT SEIZURE ACTIVITY IN PATIENTS 6 YEARS OF AGE AND OLDER
- U-2728 USE OF PEMETREXED WITH PRIOR AND/OR REPEATED VITAMIN B12 AND FOLIC ACID ADMINISTRATION IN PATIENTS WITH NON-SQUAMOUS NON-SMALL CELL LUNG CANCER
- U-2729 USE OF PEMETREXED WITH PRIOR AND/OR REPEATED VITAMIN B12 AND FOLIC ACID ADMINISTRATION IN PATIENTS WITH MESOTHELIOMA
- U-2730 METHOD OF TREATING TYPE 2 DIABETES MELLITUS USING A PHARMACEUTICAL COMPOSITION COMPRISING EMPAGLIFLOZIN, LINAGLIPTIN AND METFORMIN
- U-2731 METHOD OF TREATING TYPE 2 DIABETES MELLITUS IN A PATIENT WITH RENAL IMPAIRMENT

PATENT AND EXCLUSIVITY TERMS

ADB 167 of 225

PATENT USE

- (45 ML/MIN/1.73 M2<=EGFR<60 ML/MIN/1.73 M2) BY ONCE DAILY ADMINISTRATION OF 10 MG OR 25 MG OF EMPAGLIFLOZIN IN COMBINATION WITH LINAGLIPTIN AND METFORMIN
- U-2732 METHOD OF TREATING TYPE 2 DIABETES USING A PHARMACEUTICAL COMPOSITION COMPRISING LINAGLIPTIN, METFORMIN, EMPAGLIFLOZIN AND A BASIC AMINO ACID
- U-2733 METHOD OF TREATING A TYPE 2 DIABETES MELLITUS PATIENT WITH INSUFFICIENT GLYCEMIC CONTROL DESPITE THERAPY WITH METFORMIN USING A PHARMACEUTICAL COMPOSITION COMPRISING EMPAGLIFLOZIN, LINAGLIPTIN AND METFORMIN
- U-2734 METHOD OF TREATMENT OF IRON DEFICIENCY ANEMIA IN ADULT PATIENTS WHO HAVE INTOLERANCE TO ORAL IRON OR HAVE HAD UNSATISFACTORY RESPONSE TO ORAL IRON, WHO HAVE NON-HEMODIALYSIS DEPENDENT CHRONIC KIDNEY DISEASE, BY ADMINISTERING FERRIC DERISOMALTOSE
- U-2735 MANAGEMENT OF MODERATE TO SEVERE PAIN ASSOCIATED WITH ENDOMETRIOSIS USING 150MG ELAGOLIX WHILE CO-ADMINISTERING RIFAMPIN
- U-2736 METHOD OF TREATING EPITHELIOID SARCOMA
- U-2737 METHOD OF TREATING EPITHELIOID SARCOMA BY INHIBITING ENHANCER OF ZESTE HOMOLOG 2 (EZH2)
- U-2738 METHOD OF TREATING A LUNG METASTASIS OF EPITHELIOID SARCOMA
- U-2739 INCREASING BLOOD PRESSURE WITH AN INITIAL RATE OF ABOUT 20 NG/KG/MIN ANGIOTENSIN II IN A HUMAN SUBJECT HAVING SEPTIC SHOCK, AND TITRATING THE RATE UP.
- U-2740 INCREASING BLOOD PRESSURE WITH A RATE OF ABOUT 20 NG/KG/MIN TO ABOUT 40 NG/KG/MIN ANGIOTENSIN II IN A HUMAN SUBJECT HAVING SEPTIC SHOCK
- U-2741 TREATMENT OF CLOSTRIDIODES DIFFICILE-ASSOCIATED DIARRHEA (CDAD) IN PATIENTS FROM 6 MONTHS OF AGE AND OLDER
- U-2742 TREATMENT OF SEVERE HYPOGLYCEMIA
- U-2743 USE OF VASCEPA AS AN ADJUNCT TO STATIN THERAPY TO REDUCE THE RISK OF UNSTABLE ANGINA IN AN ADULT PATIENT WITH ESTABLISHED CARDIOVASCULAR DISEASE
- U-2744 USE OF VASCEPA AS AN ADJUNCT TO STATIN THERAPY TO REDUCE THE RISK OF STROKE IN AN ADULT PATIENT WITH ESTABLISHED CARDIOVASCULAR DISEASE
- U-2745 TREATMENT OF NEUROBLASTOMAS THAT HAVE A NEUROTROPHIC TYROSINE RECEPTOR KINASE (NTRK) GENE FUSION
- U-2746 USE OF NEXLIZET AS AN ADJUNCT TO DIET AND MAXIMALLY TOLERATED STATIN THERAPY TO LOWER LDL-C IN ADULTS WITH HETEROZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA OR ESTABLISHED ATHEROSCLEROTIC CARDIOVASCULAR DISEASE
- U-2747 USE OF NEXLETOL AS AN ADJUNCT TO DIET AND MAXIMALLY TOLERATED STATIN THERAPY TO LOWER LDL-C IN ADULTS WITH HETEROZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA OR ESTABLISHED ATHEROSCLEROTIC CARDIOVASCULAR DISEASE
- U-2748 USE OF NEXLETOL AS AN ADJUNCT TO DIET AND MAXIMALLY TOLERATED STATIN THERAPY FOR INHIBITING CHOLESTEROL SYNTHESIS TO LOWER LDL-C IN ADULTS WITH HETEROZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA OR ESTABLISHED ATHEROSCLEROTIC CARDIOVASCULAR DISEASE
- U-2749 USE OF NEXLIZET AS AN ADJUNCT TO DIET AND MAXIMALLY TOLERATED STATIN THERAPY FOR INHIBITING CHOLESTEROL SYNTHESIS TO LOWER LDL-C IN ADULTS WITH HETEROZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA OR ESTABLISHED ATHEROSCLEROTIC CARDIOVASCULAR DISEASE
- U-2750 MANAGEMENT OF MODERATE-TO-SEVERE PAIN BY INTRAVENOUS INJECTION
- U-2751 A TRANSDERMAL METHOD OF CONTRACEPTION
- U-2752 METHOD OF USING L-CYSTEINE IN AN ADMIXTURE FOR TREATING PATIENTS NEEDING PARENTERAL NUTRITION
- U-2753 INCREASING SURVIVAL IN METASTATIC CASTRATION-RESISTANT PROSTATE CANCER PATIENTS PREVIOUSLY TREATED WITH DOCETAXEL BY ADMINISTERING AS A 3 WEEK CYCLE CABAZITAXEL AFTER 5 MG DEXCHLORPHENIRAMINE, 8 MG DEXAMETHASONE, AND AN H2-AGONIST
- U-2754 TREATMENT OF POST-OPERATIVE NAUSEA AND VOMITING
- U-2755 OCULAR EXAMINATION, INTRAOCULAR PRESSURE MEASUREMENT, OR REMOVAL OF FOREIGN BODIES OR SUTURES, IN ADULT AND PEDIATRIC PATIENTS REQUIRING A DISCLOSING AGENT IN COMBINATION WITH A TOPICAL OPHTHALMIC ANESTHETIC
- U-2756 USE OF VASCEPA AS AN ADJUNCT TO STATIN THERAPY TO REDUCE THE RISK OF CARDIOVASCULAR DEATH IN AN ADULT PATIENT WITH ESTABLISHED CARDIOVASCULAR DISEASE

PATENT AND EXCLUSIVITY TERMS

ADB 168 of 225

PATENT USE

U-2757 DOSING REGIMEN FOR THE TREATMENT OF SCHIZOPHRENIA IN ADULTS BY ADMINISTERING TWO LOADING DOSES OF PALIPERIDONE PALMITATE FOLLOWED BY MAINTENANCE DOSE(S)

U-2758 DOSING REGIMEN FOR THE TREATMENT OF SCHIZOAFFECTIVE DISORDER IN ADULTS AS A MONOTHERAPY AND AS AN ADJUNCT TO MOOD STABILIZERS OR ANTIDEPRESSANTS BY ADMINISTERING TWO LOADING DOSES OF PALIPERIDONE PALMITATE FOLLOWED BY MAINTENANCE DOSE(S)

U-2759 REDUCTION OF INTRAOCULAR PRESSURE (IOP) IN PATIENTS WITH OPEN ANGLE GLAUCOMA(OAG) OR OCULAR HYPERTENSION (OHT) WITH A BIODEGRADABLE BIMATOPROST IMPLANT

U-2760 TOPICAL TREATMENT OF ACNE VULGARIS IN PATIENTS 12 YEARS OF AGE OR OLDER

U-2761 INTRAVENOUS SOTALOL DOSING REGIMEN FOR ACHIEVING STEADY STATE CONCENTRATION (EXPOSURE) FASTER COMPARED TO THE CONVENTIONAL ORAL DOSING IN A FACILITY THAT CAN PROVIDE ELECTROCARDIOGRAPHIC MONITORING

U-2762 USE OF VASCEPA AS AN ADJUNCT TO STATIN THERAPY TO REDUCE THE RISK OF A MAJOR CARDIOVASCULAR EVENT IN AN ADULT PATIENT WITH DIABETES MELLITUS AND TWO OR MORE ADDITIONAL RISK FACTORS FOR CARDIOVASCULAR DISEASE

U-2763 METHOD OF TREATING ADULTS WITH SCHIZOPHRENIA COMPRISING ADMINISTERING ASENAPINE VIA A TRANSDERMAL PATCH

U-2764 TREATMENT OF POST-OPERATIVE INFLAMMATION AND PAIN FOLLOWING OCULAR SURGERY

U-2765 TREATMENT OF HIV-1 INFECTION IN ADULTS AND PEDIATRIC PATIENTS WEIGHING AT LEAST 40 KG WHO HAVE NO PRIOR ANTIRETROVIRAL TREATMENT HISTORY OR ARE VIROLOGICALLY SUPPRESSED ON A STABLE ANTIRETROVIRAL REGIMEN FOR AT LEAST 6 MONTHS

U-2766 TX OF HIV1 INFECTION USING A COMPOSITION CONTAINING A PK ENHANCER THAT INHIBITS CY P450 MONOOXYGENASE IN ADULTS & PEDIATRIC PATIENTS AT LEAST 40KG HAVING NO PRIOR ARV TX HISTORY OR ARE VIROLOGICALLY SUPPRESSED ON A STABLE ARV REGIMEN FOR AT LEAST 6 MO

U-2767 TREATMENT OF HIV-1 INFECTION USING A COMPOSITION CONTAINING A PK ENHANCER THAT INHIBITS CY P450 MONOOXYGENASE IN ADULTS AND PEDIATRIC PATIENTS WEIGHING AT LEAST 40KG WHO HAVE NO PRIOR ARV TREATMENT HISTORY

U-2768 TREATMENT OF HIV-1 INFECTION USING A COMPOSITION CONTAINING A PK ENHANCER THAT INHIBITS CY P450 MONOOXYGENASE IN ADULTS AND PEDIATRIC PATIENTS WEIGHING AT LEAST 40KG WHO ARE VIROLOGICALLY SUPPRESSED ON A STABLE ARV REGIMEN FOR AT LEAST 6 MONTHS

U-2769 DOSING REGIMEN FOR INTRAVENOUS SOTALOL FOR ADMINISTRATION IN A FACILITY THAT CAN PROVIDE CONTINUOUS ELECTROCARDIOGRAPHIC MONITORING AND CARDIAC RESUSCITATION.

U-2770 CUSHING'S DISEASE

U-2771 TREATMENT OF MODERATE TO SEVERE PLAQUE PSORIASIS IN PATIENTS 18 YEARS OF AGE OR OLDER

U-2772 MAINTENANCE TREATMENT OF CHRONIC PULMONARY DISEASE (COPD)

U-2774 TREATMENT OF RELAPSING FORMS OF MULTIPLE SCLEROSIS (MS), TO INCLUDE CLINICALLY ISOLATED SYNDROME, RELAPSING-REMITTING DISEASE, AND ACTIVE SECONDARY PROGRESSIVE DISEASE, IN ADULTS

U-2775 TREATMENT OF A TYPE 2 DIABETES MELLITUS PATIENT WITH INSUFFICIENT GLYCEMIC CONTROL DESPITE METFORMIN THERAPY USING A COMPOSITION COMPRISING AN EXTENDED RELEASE CORE COMPRISING METFORMIN AND AN OUTER COATING COMPRISING EMPAGLIFLOZIN

U-2776 TREATMENT OF A TYPE 2 DIABETES MELLITUS PATIENT WITH INSUFFICIENT GLYCEMIC CONTROL DESPITE METFORMIN THERAPY USING A COMPOSITION COMPRISING AN EXTENDED RELEASE CORE COMPRISING METFORMIN AND AN OUTER COATING COMPRISING EMPAGLIFLOZIN AND LINAGLIPTIN

U-2777 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS 6 YEARS AND OLDER WHO ARE HOMOZYGOUS FOR THE F508DEL MUTATION IN THE CFTR GENE USING A PHARMACEUTICAL COMPOSITION ACCORDING TO CLAIM 2 OF U.S. PATENT NO. 10,597,384, FURTHER COMPRISING IVACAFTOR

U-2778 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS 2 TO 5 YEARS OLD WHO ARE HOMOZYGOUS FOR THE F508DEL MUTATION IN THE CFTR GENE USING A PHARMACEUTICAL COMPOSITION ACCORDING TO CLAIM 2 OF U.S. PATENT NO. 10,597,384, FURTHER COMPRISING IVACAFTOR

U-2779 TREATMENT OF SPASTICITY

U-2780 USE FOR THE TREATMENT OF SEIZURES IN PATIENTS WITH LENNOX-GASTAUT SYNDROME

U-2781 USE FOR THE TREATMENT OF SEIZURES IN PATIENTS WITH DRAVET SYNDROME

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-2782	USE FOR REDUCING CONVULSIVE SEIZURE FREQUENCY IN PATIENTS WITH LENNOX GASTAUT SYNDROME
U-2783	USE FOR REDUCING CONVULSIVE SEIZURE FREQUENCY IN PATIENTS WITH DRAVET SYNDROME
U-2784	A METHOD OF TREATING A CANCER CHARACTERIZED BY AN IDH1 MUTATION WHERE THE CANCER IS RELAPSED OR REFRACTORY ACUTE MYELOID LEUKEMIA (AML) AND WHERE THE MUTANT IDH1 HAS THE ABILITY TO CONVERT ALPHA-KETOGLUTARATE INTO 2-HYDROXYGLUTARATE (2-HG)
U-2785	A METHOD OF TREATING A CANCER CHARACTERIZED BY AN IDH1 MUTATION WHERE THE CANCER IS NEWLY DIAGNOSED ACUTE MYELOID LEUKEMIA (AML) AND WHERE THE MUTANT IDH1 HAS THE ABILITY TO CONVERT ALPHA-KETOGLUTARATE INTO 2-HYDROXYGLUTARATE (2-HG)
U-2786	METHOD OF PREVENTING PREGNANCY BY INSERTING A VAGINAL SYSTEM CONTAINING 103 MG OF SEGESTERONE ACETATE AND 17.4 MG ETHINYL ESTRADIOL INTO A VAGINA FOR UP TO THIRTEEN 21/7-DAY (IN/OUT) CYCLES
U-2787	METHOD OF CONTRACEPTION BY INSERTING A VAGINAL SYSTEM FOR UP TO 13 21/7-DAY (IN/OUT) CYCLES, WHEREIN EFFICACY REQUIRES THE SYSTEM CANNOT BE OUT OF THE VAGINA FOR MORE THAN 2 CUMULATIVE HOURS IN ANY SUCH CYCLE WITHOUT USING ALTERNATIVE CONTRACEPTION
U-2788	TREATMENT OF BREAST CANCER INCLUDING HER2 (ERBB2)-POSITIVE OR -OVEREXPRESSING BREAST CANCER
U-2789	POTASSIUM PHOSPHATES INJECTION IS INDICATED AS A SOURCE OF PHOSPHORUS IN INTRAVENOUS FLUIDS TO CORRECT HYPOPHOSPHATEMIA IN ADULTS AND PEDIATRIC PATIENTS 12 YEARS OF AGE AND OLDER
U-2790	TREATMENT OF A TREATMENT-NAIVE PATIENT WITH INADEQUATELY CONTROLLED TYPE 2 DIABETES USING A COMPOSITION COMPRISING AN EXTENDED RELEASE CORE COMPRISING METFORMIN AND AN OUTER COATING COMPRISING EMPAGLIFLOZIN AND LINAGLIPTIN
U-2791	TREATMENT OF ADULT PATIENTS WITH INSOMNIA, CHARACTERIZED BY DIFFICULTIES WITH SLEEP ONSET AND/OR SLEEP MAINTENANCE
U-2792	TREATMENT OF A TREATMENT-NAIVE PATIENT WITH INADEQUATELY CONTROLLED TYPE 2 DIABETES USING A COMPOSITION COMPRISING AN EXTENDED RELEASE CORE COMPRISING METFORMIN AND AN OUTER COATING COMPRISING EMPAGLIFLOZIN
U-2793	A METHOD FOR DELIVERING NITRIC OXIDE TO A PATIENT WITH PULMONARY HYPERTENSION OR HYPOXIA
U-2794	TREATMENT OF TYPE 2 DIABETES MELLITUS WITH 100 MG CANAGLIFLOZIN PER DAY
U-2795	TREATMENT OF TYPE 2 DIABETES MELLITUS WITH 300 MG CANAGLIFLOZIN PER DAY
U-2796	REDUCTION OF RISK OF MAJOR ADVERSE CARDIOVASCULAR EVENTS IN THE TREATMENT OF TYPE 2 DIABETES MELLITUS PATIENTS WITH 100 MG CANAGLIFLOZIN PER DAY
U-2797	REDUCTION OF RISK OF MAJOR ADVERSE CARDIOVASCULAR EVENTS IN THE TREATMENT OF TYPE 2 DIABETES MELLITUS PATIENTS WITH 300 MG CANAGLIFLOZIN PER DAY
U-2798	REDUCTION OF RISK OF END STAGE KIDNEY DISEASE, DOUBLING OF SERUM CREATININE, CARDIOVASCULAR DEATH, AND HOSPITALIZATION FOR HEART FAILURE IN THE TREATMENT OF TYPE 2 DIABETES MELLITUS PATIENTS WITH 100 MG CANAGLIFLOZIN PER DAY
U-2799	REDUCTION OF RISK OF END STAGE KIDNEY DISEASE, DOUBLING OF SERUM CREATININE, CARDIOVASCULAR DEATH, AND HOSPITALIZATION FOR HEART FAILURE IN THE TREATMENT OF TYPE 2 DIABETES MELLITUS PATIENTS WITH 300 MG CANAGLIFLOZIN PER DAY
U-2800	TREATMENT OF PEDIATRIC PATIENTS 2 YEARS OF AGE AND OLDER WITH NEUROFIBROMATOSIS TYPE 1 (NF1) WHO HAVE SYMPTOMATIC, INOPERABLE PLEXIFORM NEUROFIBROMAS (PN)
U-2801	METHOD OF IRON ADMINISTRATION TO TREAT PATIENTS IN NEED OF IRON REPLACEMENT THERAPY
U-2802	BRAFTOVI IS A KINASE INHIBITOR INDICATED IN COMBINATION WITH CETUXIMAB, FOR THE TREATMENT OF ADULT PATIENTS WITH METASTATIC COLORECTAL CANCER (CRC) WITH A BRAF V600E MUTATION, AS DETECTED BY AN FDA-APPROVED TEST, AFTER PRIOR THERAPY
U-2803	BRAFTOVI IS A KINASE INHIBITOR INDICATED IN COMBINATION WITH BINIMETINIB, FOR THE TREATMENT OF PATIENTS WITH UNRESECTABLE OR METASTATIC MELANOMA WITH A BRAF V600E OR V600K MUTATION, AS DETECTED BY AN FDA- APPROVED TEST
U-2804	A METHOD FOR THE IMPROVEMENT OF NEUROLOGICAL OUTCOME BY REDUCING THE INCIDENCE AND SEVERITY OF ISCHEMIC DEFICITS IN ADULT PATIENTS WITH SUBARACHNOID HEMORRHAGE (SAH) FROM RUPTURED INTRACRANIAL BERRY ANEURYSMS
U-2805	TREATMENT OF NON-24 HOUR SLEEP-WAKE DISORDER BY ADMINISTERING TASIMELTEON TO PATIENTS WITH A SMOKING HISTORY

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-2806 TREATMENT OF NON-24 HOUR SLEEP-WAKE DISORDER BY AVOIDING THE ADMINISTRATION OF FOOD

U-2807 TREATMENT OF MODERATE TO SEVERE MIGRAINE PAIN WITH PAIN FREE AT 2 HOURS POST ADMINISTRATION

U-2808 TREATMENT OF DYSKINESIA, DECREASING OFF TIME, AND INCREASING ON TIME WITHOUT TROUBLESOME DYSKINESIA IN PATIENTS WITH PARKINSON'S DISEASE RECEIVING LEVODOPA-BASED THERAPY, WITH OR WITHOUT CONCOMITANT DOPAMINERGIC MEDICATIONS

U-2809 FOR THE TREATMENT OF PREVIOUSLY TREATED, UNRESECTABLE LOCALLY ADVANCED OR METASTATIC CHOLANGIOCARCINOMA WITH A FIBROBLAST GROWTH FACTOR RECEPTOR 2 (FGFR2) FUSION OR OTHER REARRANGEMENT

U-2810 METHOD OF SUPPORTING EMBRYO IMPLANTATION AND EARLY PREGNANCY BY SUPPLEMENTATION OF CORPUS LUTEAL FUNCTION AS PART OF AN ASSISTED REPRODUCTIVE TECHNOLOGY (ART) TREATMENT PROGRAM FOR INFERTILE WOMEN

U-2811 METHOD OF TREATING PARKINSON'S DISEASE

U-2812 ADJUNCTIVE TREATMENT TO LEVODOPA/CARBIDOPA IN PATIENTS WITH PARKINSON'S DISEASE

U-2813 USE FOR THE TREATMENT OF PATIENTS WITH METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) WITH A MESENCHYMAL-EPITHELIAL TRANSITION (MET) EXON 14 SKIPPING MUTATION

U-2814 A METHOD OF PROPHYLACTIC TREATMENT OF GOUT FLARES IN ADULTS COMPRISES ADMINISTERING TO A PATIENT A LIQUID COLCHICINE ORAL SOLUTION

U-2815 MEASURING TIME-VARYING CHANGE IN BLOOD IN A TISSUE VOLUME USING MODIFIED BEER-LAMBERT LAW IN VASCULAR, GASTROINTESTINAL, ORGAN TRANSPLANT, AND PLASTIC, MICRO-AND RECONSTRUCTIVE, INCLUDING MINIMALLY INVASIVE, SURGERY

U-2816 METHOD FOR TREATING INFLUENZA

U-2817 METHOD OF INHIBITING COMT IN THE PERIPHERY

U-2818 METHOD OF REDUCING O-METHYLATION OF L-DOPA

U-2819 MAINTENANCE TREATMENT WITH BEVACIZUMAB OF ADV. EPITHELIAL OVARIAN CANCER IN COMPLETE OR PARTIAL RESPONSE TO FIRST-LINE PLATINUM CHEMOTHERAPY AND HOMOLOGOUS RECOMBINATION DEFICIENCY-POSITIVE WITH A DELETERIOUS OR SUSPECTED DELETERIOUS BRCA MUTATION

U-2820 MAINTENANCE TREATMENT WITH BEVACIZUMAB OF ADV. EPITHELIAL OVARIAN CANCER IN COMPLETE OR PARTIAL RESPONSE TO FIRST-LINE PLATINUM CHEMOTHERAPY AND HOMOLOGOUS RECOMBINATION DEFICIENCY-POSITIVE WITH GENOMIC INSTABILITY

U-2821 MAINTENANCE TREATMENT WITH BEVACIZUMAB OF FALLOPIAN TUBE CANCER IN COMPLETE OR PARTIAL RESPONSE TO FIRST-LINE PLATINUM CHEMOTHERAPY AND HOMOLOGOUS RECOMBINATION DEFICIENCY-POSITIVE WITH A DELETERIOUS OR SUSPECTED DELETERIOUS BRCA MUTATION

U-2822 MAINTENANCE TREATMENT WITH BEVACIZUMAB OF FALLOPIAN TUBE CANCER IN COMPLETE OR PARTIAL RESPONSE TO FIRST-LINE PLATINUM CHEMOTHERAPY AND HOMOLOGOUS RECOMBINATION DEFICIENCY-POSITIVE WITH GENOMIC INSTABILITY

U-2823 MAINTENANCE TREATMENT WITH BEVACIZUMAB OF PRIMARY PERITONEAL CANCER IN COMPLETE OR PARTIAL RESPONSE TO FIRST-LINE PLATINUM CHEMOTHERAPY AND HOMOLOGOUS RECOMBINATION DEFICIENCY-POSITIVE WITH A DELETERIOUS OR SUSPECTED DELETERIOUS BRCA MUTATION

U-2824 MAINTENANCE TREATMENT WITH BEVACIZUMAB OF PRIMARY PERITONEAL CANCER IN COMPLETE OR PARTIAL RESPONSE TO FIRST-LINE PLATINUM CHEMOTHERAPY AND HOMOLOGOUS RECOMBINATION DEFICIENCY-POSITIVE WITH GENOMIC INSTABILITY

U-2825 TREATMENT OF "OFF" EPISODES IN PATIENTS WITH PARKINSON'S DISEASE

U-2826 TREATMENT OF ADULT PATIENTS WITH METASTATIC RET FUSION-POSITIVE NON-SMALL CELL LUNG CANCER

U-2827 TREATMENT OF ADULT AND PEDIATRIC PATIENTS 12 YEARS OF AGE AND OLDER WITH ADVANCED OR METASTATIC RET-MUTANT MEDULLARY THYROID CANCER (MTC) WHO REQUIRE SYSTEMIC THERAPY

U-2828 TREATMENT OF ADULT AND PEDIATRIC PATIENTS 12 YEARS OF AGE AND OLDER WITH ADVANCED OR METASTATIC RET FUSION-POSITIVE THYROID CANCER WHO REQUIRE SYSTEMIC THERAPY AND WHO ARE RADIOACTIVE IODINE REFRACTORY (IF RADIOACTIVE IODINE IS APPROPRIATE)

U-2829 TREATMENT OF GASTROINTESTINAL STROMAL TUMORS

U-2830 A METHOD FOR TREATING METASTATIC CASTRATION-RESISTANT PROSTATE CANCER (MCRPC),

PATENT AND EXCLUSIVITY TERMS

ADB 171 of 225

PATENT USE

WHEREIN THE CANCER IS ASSOCIATED WITH A DELETERIOUS BRCA-MUTATION

- U-2831 TREATMENT OF PARTIAL-ONSET SEIZURES IN A PATIENT WITH REFRACTORY PARTIAL-ONSET SEIZURES
- U-2832 TREATMENT OF DELETERIOUS OR SUSPECTED DELETERIOUS GERMLINE OR SOMATIC HOMOLOGOUS RECOMBINATION REPAIR GENE-MUTATED METASTATIC CASTRATION-RESISTANT PROSTATE CANCER, WHICH HAS PROGRESSED FOLLOWING PRIOR TREATMENT WITH ENZALUTAMIDE OR ABIRATERONE
- U-2833 TREATMENT OF DELETERIOUS OR SUSPECTED DELETERIOUS GERMLINE OR SOMATIC BRCA-MUTATED METASTATIC CASTRATION-RESISTANT PROSTATE CANCER, WHICH HAS PROGRESSED FOLLOWING PRIOR TREATMENT WITH ENZALUTAMIDE OR ABIRATERONE
- U-2834 TREATMENT OF ADULTS WITH PERSISTENT/RECURRENT CHRONIC THROMBOEMBOLIC PULMONARY HYPERTENSION (CTEPH), (WHO GROUP 4) AFTER SURGICAL TREATMENT, OR INOPERABLE CTEPH, TO IMPROVE EXERCISE CAPACITY AND WHO FUNCTIONAL CLASS
- U-2835 TREATMENT OF ADULTS WITH PULMONARY HYPERTENSION (PAH), (WHO GROUP 1), TO IMPROVE EXERCISE CAPACITY, WHO FUNCTIONAL CLASS AND TO DELAY CLINICAL WORSENING
- U-2836 TREATMENT OF ADULT PATIENTS WITH SMALL CELL LUNG CANCER (SCLC) WITH DISEASE PROGRESSION ON OR AFTER PLATINUM-BASED CHEMOTHERAPY.
- U-2837 TREATMENT OF ANAPLASTIC LYMPHOMA KINASE (ALK)-POSITIVE METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC)
- U-2838 REDUCTION OF THE RATE OF A FIRST MYOCARDIAL INFARCTION OR STROKE IN PATIENTS WITH CORONARY ARTERY DISEASE AT HIGH RISK FOR SUCH EVENTS
- U-2839 TREATMENT OF MYOCARDIAL INFARCTION OR STROKE IN PATIENTS WITH CORONARY ARTERY DISEASE AT HIGH RISK FOR SUCH EVENTS
- U-2840 TREATMENT OF HOSPITAL-ACQUIRED BACTERIAL PNEUMONIA AND VENTILATOR-ASSOCIATED BACTERIAL PNEUMONIA (HABP/VABP)
- U-2841 USE OF VASCEPA WITH HIGH INTENSITY STATIN THERAPY TO REDUCE THE RISK OF A CV EVENT IN AN ADULT PATIENT WITH ELEVATED TRIGLYCERIDE LEVELS AND (1) ESTABLISHED CV DISEASE, OR (2) DIABETES MELLITUS AND TWO OR MORE ADDITIONAL RISK FACTORS FOR CV DISEASE
- U-2842 MANAGEMENT OF HEAVY MENSTRUAL BLEEDING ASSOCIATED WITH UTERINE LEIOMYOMAS (FIBROIDS)
- U-2843 NASAL ADMINISTRATION OF METOCLOPRAMIDE FOR TREATMENT OF DIABETIC GASTROPARESIS
- U-2844 IN COMBINATION WITH PEMBROLIZUMAB FOR THE FIRST-LINE TREATMENT OF PATIENTS WITH ADVANCED RENAL CELL CARCINOMA
- U-2845 A METHOD OF TREATING A HUMAN PATIENT SUFFERING FROM PULMONARY HYPERTENSION
- U-2846 TREATMENT OF CHRONIC GRAFT VERSUS HOST DISEASE REFRACTORY TO SYSTEMIC THERAPY
- U-2847 REDUCTION IN RISK OF OVERT HEPATIC ENCEPHALOPATHY (HE) RECURRENCE IN ADULTS
- U-2848 TREATMENT OF TRAVELERS' DIARRHEA (TD) CAUSED BY NONINVASIVE STRAINS OF ESCHERIA COLI IN ADULT AND PEDIATRIC PATIENTS 12 YEARS OF AGE AND OLDER
- U-2849 METHOD OF TREATING BLEPHAROPTOSIS
- U-2850 MANAGEMENT OF MODERATE TO SEVERE PAIN ASSOCIATED WITH ENDOMETRIOSIS USING 150 MG ELAGOLIX WHILE CO-ADMINISTERING KETOCONAZOLE
- U-2851 METHOD OF TREATING RELAPSED OR REFRACTORY FOLLICULAR LYMPHOMA POSITIVE FOR AN ENHANCER OF ZESTE HOMOLOG 2 (EZH2) MUTATION BY INHIBITING EZH2
- U-2852 METHOD OF TREATING RELAPSED OR REFRACTORY FOLLICULAR LYMPHOMA POSITIVE FOR AN ENHANCER OF ZESTE HOMOLOG 2 (EZH2) MUTATION
- U-2853 METHOD OF TREATING RELAPSED OR REFRACTORY FOLLICULAR LYMPHOMA
- U-2854 METHOD OF TREATING RELAPSED OR REFRACTORY FOLLICULAR LYMPHOMA BY INHIBITING EZH2
- U-2855 XPOVIO IS INDICATED FOR THE TREATMENT OF ADULT PATIENTS WITH RELAPSED OR REFRACTORY DIFFUSE LARGE B-CELL LYMPHOMA (DLBCL), NOT OTHERWISE SPECIFIED, INCLUDING DLBCL ARISING FROM FOLLICULAR LYMPHOMA, AFTER AT LEAST 2 LINES OF SYSTEMIC THERAPY
- U-2856 INCREASING SURVIVAL IN METASTATIC CASTRATION-RESISTANT PROSTATE CANCER PATIENTS PREVIOUSLY TREATED WITH DOCETAXEL BY ADMINISTERING 20 TO 25 MG/M2 CABAZITAXEL AFTER A PREMEDICATION REGIMEN THAT INCLUDES AN H2-ANTAGONIST
- U-2857 USE OF ORAL OCTREOTIDE FOR LONG-TERM MAINTENANCE TREATMENT IN ACROMEGALY PATIENTS WHO HAVE RESPONDED TO AND TOLERATED TREATMENT WITH OCTREOTIDE OR

PATENT AND EXCLUSIVITY TERMS

ADB 172 of 225

PATENT USE

LANREOTIDE

- U-2858 USE IN COMBINATION WITH STIRIPENTOL, VALPROATE, AND CLOBAZAM FOR THE TREATMENT OF SEIZURES ASSOCIATED WITH DRAVET SYNDROME
- U-2859 USE OF CARDIAC MONITORING AND RESTRICTED DISTRIBUTION OF FENFLURAMINE TO MITIGATE RISK OF CARDIOVASCULAR TOXICITY IN THE TREATMENT OF SEIZURES ASSOCIATED WITH DRAVET SYNDROME
- U-2860 USE IN COMBINATION WITH STIRIPENTOL FOR THE TREATMENT OF SEIZURES ASSOCIATED WITH DRAVET SYNDROME
- U-2861 USE IN COMBINATION WITH CANNABIDIOL FOR THE TREATMENT OF SEIZURES ASSOCIATED WITH DRAVET SYNDROME
- U-2862 USE FOR THE TREATMENT OF FOCAL SEIZURES IN PATIENTS WITH DRAVET SYNDROME
- U-2863 TOPICAL TREATMENT OF HEAD LICE INFESTATION IN PATIENTS 6 MONTHS OF AGE AND OLDER
- U-2864 METHOD FOR INHIBITING CYTIDINE DEAMINASE BY ADMINISTERING CEDAZURIDINE
- U-2865 TREATMENT OF MYELODYSPLASTIC SYNDROME
- U-2866 TREATMENT OF CHRONIC MYELOMONOCYTIC LEUKEMIA
- U-2867 METHOD FOR INHIBITING DEGRADATION OF A CDA SUBSTRATE BY ADMINISTERING CEDAZURIDINE
- U-2868 TREATMENT OF NON-INFECTIOUS UVEITIS AFFECTING THE POSTERIOR SEGMENT OF THE EYE
- U-2869 IV ADMINISTRATION OF CANGRELOR BEFORE PCI AND CONTINUOUS INFUSION FOR AT LEAST 2 HOURS OR THE DURATION OF THE PCI AND, DURING OR AFTER THE CONTINUOUS INFUSION, ADMINISTRATION OF A LOADING DOSE OF TICAGRELOR OR AN EQUIVALENT THERAPY (PER LABELING)
- U-2870 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETIS BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 1 AND 2
- U-2871 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETIS BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 1 AND 2 AND 3
- U-2872 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETIS BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 4 AND 5
- U-2873 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETIS BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 4, 5, AND 6
- U-2874 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETIS BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING MICRONIZED BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 7 AND 8
- U-2875 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETIS BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING MICRONIZED BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 7, 8, AND 9
- U-2876 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETIS BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING MICRONIZED BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 10 AND 11
- U-2877 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETIS BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING MICRONIZED BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 10, 11, AND 12
- U-2878 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETIS BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING MICRONIZED BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 13 AND 14
- U-2879 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETIS BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING MICRONIZED BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 13, 14, AND 15
- U-2880 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETIS BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING MICRONIZED BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 16 AND 17
- U-2881 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETIS BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING MICRONIZED BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 16, 17, AND 18

PATENT AND EXCLUSIVITY TERMS

ADB 173 of 225

PATENT USE

- U-2882 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETIS BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING MICRONIZED BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 19 AND 20
- U-2883 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETIS BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING MICRONIZED BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 19, 20, AND 21
- U-2884 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETIS BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING MICRONIZED BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 22, 23, AND 24
- U-2885 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETIS BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING MICRONIZED BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 25 AND 26
- U-2886 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETIS BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING MICRONIZED BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 25, 26, AND 27
- U-2887 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETIS BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING MICRONIZED BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 28 AND 29
- U-2888 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETIS BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING MICRONIZED BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 28, 29, AND 30
- U-2889 USE FOR THE MAINTENANCE TREATMENT OF PATIENTS WITH CHRONIC OBSTRUCTIVE PULMONARY DISEASE (COPD)
- U-2890 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIMS 1 AND 2 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 1
- U-2891 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIMS 1 AND 3 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 1
- U-2892 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIMS 1 AND 4 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 1
- U-2893 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIMS 1 AND 5 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 1
- U-2894 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIMS 1 AND 6 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 1
- U-2895 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIMS 1 AND 7 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 1
- U-2896 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIMS 1 AND 8 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 1
- U-2897 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIM 1 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIMS 1 AND 9
- U-2898 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIM 1 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIMS 1 AND 10
- U-2899 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIM 1 AND WHEREIN THE EFFECTS ARE AS RECITED IN

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

CLAIMS 1 AND 11

U-2900	ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIM 1 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIMS 1 AND 13
U-2901	ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIMS 1 AND 15 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 1
U-2902	ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIMS 1 AND 16 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 1
U-2903	ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIMS 1 AND 17 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 1
U-2904	ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIMS 1, 17 AND 18 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 1
U-2905	ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIMS 1 AND 19 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 1
U-2906	ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIM 22 WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 22
U-2907	ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIMS 22 AND 23 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 22
U-2908	ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIMS 22 AND 24 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 22
U-2909	ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIMS 22 AND 25 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 22
U-2910	ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIM 26 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 26
U-2911	ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIMS 26 AND 27 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 26
U-2912	ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIMS 26 AND 28 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 26
U-2913	ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIMS 26 AND 29 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 26
U-2914	ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIMS 26 AND 30 WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 26
U-2915	ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2

PATENT AND EXCLUSIVITY TERMS

ADB 175 of 225

PATENT USE

- DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIMS 26 AND 31 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 26
- U-2916 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIMS 26 AND 32 WHEREIN THE EFFECTS ARE AS RECITED IN SAID CLAIMS
- U-2917 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIMS 26 AND 33 WHEREIN THE EFFECTS ARE AS RECITED IN CLAIMS
- U-2918 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIMS 26 AND 34 WHEREIN THE EFFECTS ARE AS RECITED IN SAID CLAIMS
- U-2919 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIMS 26 AND 36 WHEREIN THE EFFECTS ARE AS RECITED IN SAID CLAIMS
- U-2920 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIMS 26 AND 38 AND WHEREIN THE EFFECTS ARE AS RECITED CLAIM 26
- U-2921 ONCE DAILY TOPICAL TREATMENT OF PERSISTENT FACIAL ERYTHEMA ASSOCIATED WITH ROSACEA IN ADULTS
- U-2922 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIMS 26 AND 39 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 26
- U-2923 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIMS 26 AND 40 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 26
- U-2924 ADJUNCT TO DIET AND EXERCISE TO TREAT GLUCOSE INTOLERANCE IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIM 42 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 42
- U-2925 ADJUNCT TO DIET AND EXERCISE TO TREAT GLUCOSE INTOLERANCE IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIMS 42 AND 43 WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 42
- U-2926 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIM 44 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 44
- U-2927 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIMS 44 AND 45 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 44
- U-2928 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIMS 44 AND 46 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 44
- U-2929 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIMS 44 AND 47 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 44
- U-2930 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIMS 44 AND 48 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 44
- U-2931 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2

PATENT AND EXCLUSIVITY TERMS

ADB 176 of 225

PATENT USE

- DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIMS 44 AND 49 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 44
- U-2932 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIMS 44 AND 50 WHEREIN THE EFFECTS ARE AS RECITED IN SAID CLAIMS
- U-2933 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIMS 44 AND 51 WHEREIN THE EFFECTS ARE AS RECITED IN SAID CLAIMS
- U-2934 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIMS 44 AND 52 WHEREIN THE EFFECTS ARE AS RECITED IN SAID CLAIMS
- U-2935 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIM 44 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIMS 44 AND 54
- U-2936 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIMS 44 AND 56 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 44
- U-2937 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIMS 44 AND 57 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 44
- U-2938 TREATMENT OF CATAPLEXY OR EXCESSIVE DAYTIME SLEEPINESS (EDS) IN PATIENTS 7 YEARS OF AGE AND OLDER WITH NARCOLEPSY WITH A MIXTURE OF SODIUM, POTASSIUM, MAGNESIUM, AND CALCIUM SALTS OF GHB
- U-2939 TREATMENT OF HIV INFECTION IN ADULTS AND PEDIATRIC PATIENTS WEIGHING AT LEAST 40KG USING A COMPOSITION CONTAINING A PHARMACOKINETIC ENHANCER THAT INHIBITS CYTOCHROME P450 MONOOXYGENASE
- U-2940 METHOD OF TREATING PEDIATRIC PATIENTS 2 YEARS OF AGE AND OLDER WITH CENTRAL PRECOCIOUS PUBERTY
- U-2941 A METHOD OF USING AN AEROSOL DELIVERY DEVICE TO AEROSOLIZE GLYCOPYRROLATE FOR THE TREATMENT OF CHRONIC OBSTRUCTIVE PULMONARY DISEASE (COPD)
- U-2942 METHOD OF TREATING ACNE VULGARIS WITH TOPICALLY APPLIED CORTEXOLONE 17A-PROPIONATE
- U-2943 TREATMENT OF RELAPSED OR REFRACTORY CHRONIC LYMPHOCYTIC LEUKEMIA
- U-2944 TREATMENT OF RELAPSED OR REFRACTORY SMALL LYMPHOCYTIC LYMPHOMA
- U-2945 TREATMENT OF PRIMARY BILIARY CHOLANGITIS (PBC) IN COMBINATION WITH URSODEOXYCHOLIC ACID (UDCA) IN ADULTS WITH AN INADEQUATE RESPONSE TO UDCA, OR AS MONOTHERAPY IN ADULTS UNABLE TO TOLERATE UDCA
- U-2946 TREATMENT OF COLORECTAL CANCER THAT HAS A NEUROTROPHIC TYROSINE RECEPTOR KINASE (NTRK) GENE FUSION
- U-2947 KYPROLIS IS INDICATED IN COMBINATION WITH DARATUMUMAB PLUS DEXAMETHASONE FOR THE TREATMENT OF PATIENTS WITH RELAPSED OR REFRACTORY MULTIPLE MYELOMA WHO HAVE RECEIVED ONE TO THREE LINES OF THERAPY
- U-2948 A METHOD OF POSITIONING AN INTRAUTERINE SYSTEM (IUS) BY DETERMINING A DEPTH OF THE UTERUS, HOLDING AN INSERTER HANDLE WITH ONE HAND, INSERTING THE IUS INTO THE UTERUS, AND RETRACTING A SLIDER ON THE HANDLE TO RELEASE THE IUS INTO THE UTERUS
- U-2949 A METHOD FOR INDUCING A POST-SURGICAL ANALGESIA SPARING EFFECT BY IMPLANTING AT THE SURGICAL SITE A COLLAGEN SPONGE CONTAINING BUPIVACAINE HCL WHICH PROVIDES LOCAL ANESTHESIA FOR UP TO 24 HOURS FOLLOWING IMPLANTATION
- U-2950 CONTINUED TREATMENT OF ADULTS WITH ACUTE MYELOID LEUKEMIA WHO ACHIEVED FIRST COMPLETE REMISSION (CR) OR CR WITH INCOMPLETE BLOOD COUNT RECOVERY FOLLOWING INTENSIVE INDUCTION CHEMOTHERAPY AND ARE NOT ABLE TO COMPLETE INTENSIVE CURATIVE THERAPY
- U-2951 USE OF CU-64 DOTATATE WITH POSITRON EMISSION TOMOGRAPHY (PET) FOR LOCALIZATION

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

- OF SOMATOSTATIN RECEPTOR POSITIVE NEUROENDOCRINE TUMORS (NETS) IN ADULT PATIENTS
- U-2952 TREATMENT OF ADULT PATIENTS WITH METASTATIC REARRANGED DURING TRANSFECTION (RET) FUSION-POSITIVE NON-SMALL CELL LUNG CANCER (NSCLC) AS DETECTED BY AN FDA APPROVED TEST
- U-2953 USE OF FLUTICASONE FUROATE FOR THE TREATMENT OF AN INFLAMMATORY OR ALLERGIC CONDITIONS, INCLUDING CHRONIC OBSTRUCTIVE PULMONARY DISEASE AND ASTHMA
- U-2954 METHOD OF DISPENSING A COMBINATION MEDICAMENT PRODUCT FROM CLAIMED DELIVERY DEVICE, FOR EXAMPLE FOR THE TREATMENT OF ASTHMA OR COPD
- U-2955 INDICATED FOR THE LONG-TERM, ONCE-DAILY, MAINTENANCE TREATMENT OF PATIENTS WITH CHRONIC OBSTRUCTIVE PULMONARY DISEASE (COPD), INCLUDING CHRONIC BRONCHITIS AND/OR EMPHYSEMA; AND ASTHMA
- U-2956 METHOD OF TREATING LAMBERT-EATON MYASTHENIC SYNDROME WITH AMIFAMPRIDINE
- U-2957 MAINTENANCE TREATMENT OF ASTHMA IN PATIENTS 18 YRS AND OLDER, OR CHRONIC OBSTRUCTIVE PULMONARY DISEASE (COPD) INCLUDING CHRONIC BRONCHITIS AND/OR EMPHYSEMA
- U-2958 A METHOD FOR CONTRACEPTION, THE METHOD COMPRISING ADMINISTERING A TABLET COMPRISING 20 MG TO 30 MG OF ULIPRISTAL ACETATE TO A WOMAN WITHIN 120 HOURS AFTER AN UNPROTECTED INTERCOURSE
- U-2959 USE OF VASCEPA AS AN ADJUNCT TO STATIN THERAPY TO REDUCE THE RISK OF A THIRD AND FURTHER CARDIOVASCULAR EVENT IN AN ADULT PATIENT WITH ELEVATED TG LEVELS (≥ 150 MG/DL) AND ESTABLISHED CARDIOVASCULAR DISEASE
- U-2960 USE OF VASCEPA AS AN ADJUNCT TO STATIN THERAPY TO REDUCE THE RISK OF A SECOND OR FURTHER CARDIOVASCULAR (CV) EVENT IN AN ADULT PATIENT WITH ELEVATED TG LEVELS (≥ 150 MG/DL) AND DIABETES MELLITUS AND 2 OR MORE ADDITIONAL RISK FACTORS FOR CV DISEASE
- U-2961 USE OF VASCEPA AS AN ADJUNCT TO STATIN THERAPY TO REDUCE THE RISK OF MYOCARDIAL INFARCTION, STROKE, BOTH IN AN ADULT PATIENT WITH TYPE 2 DIABETES MELLITUS
- U-2962 USE OF VASCEPA AS AN ADJUNCT TO STATIN THERAPY TO REDUCE THE RISK OF CORONARY REVASCULARIZATION IN AN ADULT PATIENT WITH ESTABLISHED CARDIOVASCULAR DISEASE
- U-2963 TREATMENT OF CYSTIC FIBROSIS USING IVACAFTOR IN A PATIENT AGE 4 MONTHS TO <6 YEARS WHO HAS ONE MUTATION IN THE CFTR GENE THAT IS RESPONSIVE TO IVACAFTOR BASED ON CLINICAL AND/OR IN VITRO ASSAY DATA
- U-2964 TREATMENT OF CYSTIC FIBROSIS USING IVACAFTOR IN A PATIENT AGE 4 MONTHS TO <6 YEARS WHO HAS A R117H MUTATION IN THE CFTR GENE
- U-2965 TREATMENT OF A MODERATE TO MILD CLINICAL PHENOTYPE OF CF USING IVACAFTOR IN A PATIENT AGE 4 MONTHS TO <6 YEARS WHO HAS ONE CFTR MUTATION RESPONSIVE TO IVACAFTOR BASED ON CLINICAL AND/OR IN VITRO ASSAY DATA
- U-2966 TREATMENT OF CF IN A PATIENT AGE 4 MONTHS TO <6 YEARS WHO HAS ONE CFTR MUTATION RESPONSIVE TO IVACAFTOR BASED ON CLINICAL AND/OR IN VITRO ASSAY DATA USING A SOLID COMPOSITION COMPRISING AMORPHOUS (LESS THAN ABOUT 30% CRYSTALLINE) IVACAFTOR
- U-2967 TREATMENT OF CF IN A PATIENT AGE 4 MONTHS TO <6 YEARS WHO HAS ONE MUTATION IN THE CFTR GENE THAT IS RESPONSIVE TO IVACAFTOR BASED ON CLINICAL AND/OR IN VITRO ASSAY DATA USING THE COMPOSITION RECITED IN CLAIM 1 OF US 10272046
- U-2968 USE OF REMIMAZOLAM FOR INDUCTION AND MAINTENANCE OF PROCEDURAL SEDATION IN ADULTS UNDERGOING PROCEDURES LASTING 30 MINUTES OR LESS
- U-2969 TREATMENT OF ADULT PATIENTS WITH CYCLOSPORIN-RESISTANT, STEROID-DEPENDENT/REFRACTORY, OR STEROID RESISTANT CHRONIC GRAFT-VERSUS-HOST DISEASE (CGVHD) AFTER FAILURE OF ONE OR MORE LINES OF SYSTEMIC THERAPY
- U-2970 TREATMENT OF ADULT PATIENTS WITH CHRONIC GRAFT-VERSUS-HOST DISEASE (CGVHD) AFTER FAILURE OF ONE OR MORE LINES OF SYSTEMIC THERAPY
- U-2971 THE TREATMENT OF ADULT PATIENTS WITH METASTATIC RET FUSION-POSITIVE NON-SMALL CELL LUNG CANCER (NSCLC)
- U-2972 THE TREATMENT OF ADULT AND PEDIATRIC PATIENTS 12 YEARS OF AGE AND OLDER WITH ADVANCED OR METASTATIC RET-MUTANT MEDULLARY THYROID CANCER (MTC) WHO REQUIRE SYSTEMIC THERAPY
- U-2973 TREATING ADULT AND PEDIATRIC PATIENTS 12 YEARS OF AGE AND OLDER WITH ADVANCED OR METASTATIC RET FUSION-POSITIVE THYROID CANCER WHO REQUIRE SYSTEMIC THERAPY AND WHO ARE RADIOACTIVE IODINE-REFRACTORY, WHEREIN THE CANCER IS PAPILLARY THYROID CANCER

PATENT AND EXCLUSIVITY TERMS

ADB 178 of 225

PATENT USE

- U-2974 TREATING ADULT AND PEDIATRIC PATIENTS 12 YEARS OF AGE AND OLDER WITH ADVANCED OR METASTATIC RET FUSION-POSITIVE THYROID CANCER WHO REQUIRE SYSTEMIC THERAPY AND WHO ARE RADIOACTIVE IODINE-REFRACTORY, WHEREIN THE CANCER IS MEDULLARY THYROID CANCER
- U-2975 TREATING ADULT AND PEDIATRIC PATIENTS 12 YEARS OF AGE AND OLDER WITH ADVANCED OR METASTATIC RET FUSION-POSITIVE THYROID CANCER WHO REQUIRE SYSTEMIC THERAPY AND WHO ARE RADIOACTIVE IODINE-REFRACTORY, WHEREIN THE CANCER IS DIFFERENTIATED THYROID CANCER
- U-2976 TREATING ADULT AND PEDIATRIC PATIENTS 12 YEARS OF AGE AND OLDER WITH ADVANCED OR METASTATIC RET FUSION-POSITIVE THYROID CANCER WHO REQUIRE SYSTEMIC THERAPY AND WHO ARE RADIOACTIVE IODINE-REFRACTORY, WHEREIN THE CANCER IS RECURRENT THYROID CANCER
- U-2977 TREATING ADULT AND PEDIATRIC PATIENTS 12 YEARS OF AGE AND OLDER WITH ADVANCED OR METASTATIC RET FUSION-POSITIVE REFRACTORY DIFFERENTIATED THYROID CANCER WHO REQUIRE SYSTEMIC THERAPY AND ARE RADIOACTIVE IODINE-REFRACTORY
- U-2978 TREATMENT OF HIV-1 INFECTION IN ADULT OR PEDIATRIC PATIENTS (≥ 40 KG) WITH < 50 COPIES/ML HIV-1 RNA AFTER ≥ 6 MONTHS ON PRIOR ANTIRETROVIRAL REGIMEN AND NO KNOWN DARUNAVIR OR TENOFOVIR RESISTANCE-ASSOCIATED SUBSTITUTIONS
- U-2979 METHOD COMPRISING IV ADMINISTRATION OF CANGRELOR BEFORE PCI THEN CONTINUOUS INFUSION FOR AT LEAST 2 HOURS OR THE DURATION OF PCI AND, DURING OR AFTER CONTINUOUS INFUSION, ADMINISTRATION OF A LOADING DOSE OF TICAGRELOR, OR AN EQUIVALENT METHOD
- U-2980 METHOD OF TREATING AN ALLERGIC REACTION USING AN AUTO-INJECTOR
- U-2981 A METHOD OF TREATING ACUTE MYELOGENOUS LEUKEMIA (AML) IN A SUBJECT BY ADMINISTERING A PHARMACEUTICAL COMPOSITION WHERE THE AML IS CHARACTERIZED BY THE PRESENCE OF A MUTANT ALLELE OF IDH1 AND THE AML IS NEWLY DIAGNOSED
- U-2982 A METHOD OF TREATING ACUTE MYELOGENOUS LEUKEMIA (AML) IN A SUBJECT BY ADMINISTERING A PHARMACEUTICAL COMPOSITION WHERE THE AML IS CHARACTERIZED BY THE PRESENCE OF A MUTANT ALLELE OF IDH1 AND WHERE THE AML IS RELAPSED/REFRACTORY
- U-2983 TREATMENT OF SCHIZOPHRENIA BY RAPID AND CONTINUOUS INJECTION
- U-2984 TREATMENT OF CORONAVIRUS DISEASE 2019 (COVID-19) IN ADULTS AND PEDIATRIC PATIENTS (AT LEAST 12 YEARS OF AGE AND 40 KG) REQUIRING HOSPITALIZATION
- U-2985 A METHOD FOR TREATING DRY EYE IN A PATIENT
- U-2986 MANAGEMENT OF ACUTE PAIN BY INTRAVENOUS INJECTION
- U-2987 METHOD OF TREATING LUNG CANCER, UNDIFFERENTIATED SARCOMA, OR COLORECTAL CANCER THAT EXHIBITS AN NTRK GENE FUSION
- U-2988 REDUCTION OF THE RISK OF STROKE IN PATIENTS WITH ACUTE ISCHEMIC STROKE OR HIGH-RISK TRANSIENT ISCHEMIC ATTACK
- U-2989 METHOD OF USE AS ADJUNCTIVE THERAPY IN THE TREATMENT OF PRIMARY GENERALIZED TONIC-CLONIC SEIZURES IN PATIENTS 4 YEARS OF AGE AND OLDER
- U-2990 METHOD OF USE FOR TREATMENT OF PARTIAL-ONSET SEIZURES IN PATIENTS 4 YEARS OF AGE AND OLDER
- U-2991 METHOD OF TREATING PULMONARY ARTERIAL HYPERTENSION COMPRISING ADMINISTERING A TABLET CONTAINING SELEXIPAG
- U-2992 METHOD OF TREATING PULMONARY ARTERIAL HYPERTENSION COMPRISING ADMINISTERING A SOLID PREPARATION CONTAINING SELEXIPAG
- U-2993 A METHOD FOR TREATING A SUBJECT HAVING ADHD, SAID METHOD COMPRISING ORALLY ADMINISTERING TO SAID SUBJECT A RACEMIC METHYLPHENIDATE CHEWABLE TABLET AS CLAIMED
- U-2994 REDUCTION OF RISK OF MYOCARDIAL INFARCTION, STROKE OR CARDIOVASCULAR DEATH IN A PATIENT WITH CHRONIC CAD OR PAD BY ADMINISTERING CLINICALLY PROVEN EFFECTIVE AMOUNTS THAT ARE 2.5 MG RIVAROXABAN TWICE DAILY AND 75-100 MG ASPIRIN DAILY
- U-2995 TREATMENT OF PRIMARY HYPEROXALURIA TYPE 1 (PH1)
- U-2996 ADMINISTRATION OF AN EXTENDED RELEASE TABLET FOR THE TREATMENT OF OVERACTIVE BLADDER (OAB) WITH SYMPTOMS OF URGE URINARY INCONTINENCE, URGENCY, AND URINARY FREQUENCY
- U-2997 USE OF VASCEPA AS AN ADJUNCT TO STATIN THERAPY TO REDUCE THE RISK OF STROKE IN AN ADULT PATIENT WITH ELEVATED TRIGLYCERIDES AND ATRIAL FIBRILLATION

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

- U-2998 METHOD OF ADMINISTERING DOCETAXEL TO A SUBJECT COMBINING THE DOCETAXEL PRO-EMULSION FORMULATION WITH AN AQUEOUS MEDIUM TO PRODUCE DOCETAXEL EMULSION
- U-2999 METHOD OF USE OF TREATING, AS AN INITIAL LOADING DOSE FOR MONOTHERAPY OR ADJUNCTIVE THERAPY IN PARTIAL ONSET SEIZURE PATIENTS WITH EPILEPSY AGED 17 YEARS OR OLDER
- U-3000 METHOD FOR POST-EXPOSURE PROPHYLAXIS OF INFLUENZA
- U-3001 PROCEDURES IN ADULT AND PEDIATRIC PATIENTS REQUIRING A DISCLOSING AGENT IN COMBINATION WITH A TOPICAL OPHTHALMIC ANESTHETIC.
- U-3002 TREATMENT OF ADULT AND PEDIATRIC PATIENTS 12 YEARS OF AGE AND OLDER WITH ADVANCED OR METASTATIC RET FUSION-POSITIVE THYROID CANCER WHO REQUIRE SYSTEMIC THERAPY AND WHO ARE RADIOACTIVE IODINE-REFRACTORY (IF RADIOACTIVE IODINE IS APPROPRIATE)
- U-3003 TREATMENT OF NIGHTTIME SLEEP DISTURBANCES IN SMITH-MAGENIS SYNDROME BY ADMINISTERING TASIMELTEON
- U-3004 TREATMENT OF NIGHTTIME SLEEP DISTURBANCES IN SMITH-MAGENIS SYNDROME BY AVOIDING THE AVOIDING THE USE OF TASIMELTEON IN COMBINATION WITH A STRONG CYP1A2 INHIBITOR
- U-3005 TREATMENT OF NIGHTTIME SLEEP DISTURBANCES IN SMITH-MAGENIS SYNDROME BY AVOIDING THE USE OF TASIMELTEON WITH RIFAMPIN
- U-3006 TREATMENT OF NIGHTTIME SLEEP DISTURBANCES IN SMITH-MAGENIS SYNDROME NON-24 HOUR SLEEP-WAKE DISORDER BY AVOIDING THE USE OF TASIMELTEON IN COMBINATION WITH CYP1A2 STRONG INHIBITORS
- U-3007 TREATMENT OF NIGHTTIME SLEEP DISTURBANCES IN SMITH-MAGENIS SYNDROME BY AVOIDING THE ADMINISTRATION OF TASIMELTEON WITH FOOD
- U-3008 TREATMENT OF NIGHTTIME SLEEP DISTURBANCES IN SMITH-MAGENIS SYNDROME BY ADMINISTERING TASIMELTEON TO PATIENTS WITH A SMOKING HISTORY
- U-3009 TREATMENT OF NIGHTTIME SLEEP DISTURBANCES IN SMITH-MAGENIS SYNDROME BY ADMINISTERING TASIMELTEON TO PATIENTS WITH A SMOKING HISTORY
- U-3010 PROPHYLAXIS TO PREVENT ATTACKS OF HEREDITARY ANGIOEDEMA (HAE) IN ADULTS AND PEDIATRIC PATIENTS 12 YEARS OF AGE AND OLDER
- U-3011 A METHOD FOR TREATING OCULAR INFLAMMATION
- U-3012 TREATMENT IN COMBINATION WITH ANDROGEN DEPRIVATION THERAPY OF NON-METASTATIC, CASTRATION-RESISTANT PROSTATE CANCER (NMCRPC) THAT IMPROVES METASTASIS FREE SURVIVAL
- U-3013 TREATMENT IN COMBINATION WITH ORCHIECTOMY OF NON-METASTATIC, CASTRATION-RESISTANT PROSTATE CANCER (NMCRPC)
- U-3014 METHOD FOR THE INDUCTION OF LOCAL ANESTHESIA OF THE MUCOUS MEMBRANES
- U-3015 TOPICAL TREATMENT OF ACTINIC KERATOSIS OF THE FACE OR SCALP
- U-3016 ADJUVANT THERAPY AFTER TUMOR RESECTION IN PATIENTS WITH NON-SMALL CELL LUNG CANCER (NSCLC) WHOSE TUMORS HAVE EPIDERMAL GROWTH FACTOR RECEPTOR (EGFR) EXON 19 DELETIONS OR EXON 21 L858R MUTATIONS
- U-3017 METHOD OF TREATING EXCESSIVE DAYTIME SLEEPINESS AND/OR CATAPLEXY IN NARCOLEPSY PATIENTS WITH A SALT OF GAMMA-HYDROXYBUTYRATE WHEN DIVALPROEX SODIUM IS CONCOMITANTLY ADMINISTERED
- U-3018 XPOVIO IS INDICATED IN COMBINATION WITH BORTEZOMIB AND DEXAMETHASONE FOR THE TREATMENT OF ADULT PATIENTS WITH MULTIPLE MYELOMA WHO HAVE RECEIVED AT LEAST ONE PRIOR THERAPY
- U-3019 TREATMENT OF ADULT PATIENTS WITH ADVANCED PROSTATE CANCER THAT IS SEX-HORMONE-DEPENDENT
- U-3020 TREATMENT OF ADULT PATIENTS WITH ADVANCED PROSTATE CANCER
- U-3021 TEZACAFTOR AND IVACAFTOR FOR THE TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGE 6 YEARS AND OLDER WHO HAVE A R117H MUTATION IN THE CFTR GENE
- U-3022 TREATMENT OF CF IN PATIENTS 12 YEARS AND OLDER WHO HAVE A F508DEL OR G551D CFTR MUTATION AND A 2ND MUTATION SELECTED FROM R117H, A455E, 2789+5G->A, & 3849+10KBC->T, COMPRISING CONCURRENT COADMINISTRATION OF THE COMPOSITIONS OF CLAIM 1 OF US 10058546
- U-3023 TREATMENT OF CF IN PATIENTS 6 YEARS AND OLDER WHO HAVE A F508DEL OR G551D CFTR MUTATION AND A 2ND MUTATION SELECTED FROM R117H, A455E, 2789+5G->A, AND

PATENT AND EXCLUSIVITY TERMS

ADB 180 of 225

PATENT USE

- 3849+10KBC->T, COMPRISING CONCURRENT COADMINISTRATION OF THE COMPOSITIONS OF CLAIM 1 OF US 10058546
- U-3024 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGE 12 AND OLDER, WHO HAVE TWO COPIES OF THE F508DEL MUTATION OR WHO HAVE AT LEAST ONE CFTR MUTATION THAT IS RESPONSIVE TO TEZACAFTOR/IVACAFTOR, WITH THE COMPOSITION OF CLAIM 1 OF US 10,081,621
- U-3025 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGE 6 AND OLDER, WHO HAVE TWO COPIES OF THE F508DEL MUTATION OR WHO HAVE AT LEAST ONE CFTR MUTATION THAT IS RESPONSIVE TO TEZACAFTOR/IVACAFTOR, WITH THE COMPOSITION OF CLAIM 1 OF US 10,081,621
- U-3026 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGE 12 AND OLDER, WHO HAVE TWO COPIES OF THE F508DEL MUTATION OR AT LEAST ONE CFTR MUTATION THAT IS RESPONSIVE TO TEZACAFTOR/IVACAFTOR, WITH THE COMPOSITION OF CLAIM 1 OF US 10,206,877
- U-3027 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGE 6 AND OLDER, WHO HAVE TWO COPIES OF THE F508DEL MUTATION OR AT LEAST ONE CFTR MUTATION THAT IS RESPONSIVE TO TEZACAFTOR/IVACAFTOR, WITH THE COMPOSITION OF CLAIM 1 OF US 10,206,877
- U-3028 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 12 YEARS AND OLDER WHO HAVE AT LEAST ONE F508DEL MUTATION IN THE CFTR GENE OR A MUTATION IN THE CFTR GENE THAT IS RESPONSIVE BASED ON IN VITRO DATA WITH ELEXACAFTOR, TEZACAFTOR, AND IVACAFTOR
- U-3029 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 12 YEARS AND OLDER WHO HAVE A R117H MUTATION IN THE CFTR GENE WITH ELEXACAFTOR, TEZACAFTOR, AND IVACAFTOR
- U-3030 TREATMENT OF A MODERATE TO MILD CLINICAL PHENOTYPE OF CF IN PATIENTS AGED 12 YEARS AND OLDER WHO HAVE AT LEAST ONE F508DEL MUTATION IN THE CFTR GENE OR A MUTATION IN THE CFTR GENE THAT IS RESPONSIVE BASED ON IN VITRO DATA WITH ELX, TEZ, AND IVA
- U-3031 TREATMENT OF CF IN PATIENTS 12 YEARS AND OLDER WHO HAVE AT LEAST ONE F508DEL MUTATION OR A MUTATION IN THE CFTR GENE THAT IS RESPONSIVE BASED ON IN VITRO DATA USING A SOLID COMPOSITION COMPRISING ELX, TEZ, AMORPHOUS IVA, AND < ~30% CRYSTALLINE IVA
- U-3032 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 12 YEARS AND OLDER WHO HAVE AT LEAST ONE F508DEL MUTATION IN THE CFTR GENE OR A MUTATION IN THE CFTR GENE THAT IS RESPONSIVE BASED ON IN VITRO DATA WITH A COMPOSITION ACCORDING TO CLAIM 1 OF US 10081621
- U-3033 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 12 YEARS AND OLDER WHO HAVE AT LEAST ONE F508DEL MUTATION IN THE CFTR GENE OR A MUTATION IN THE CFTR GENE THAT IS RESPONSIVE BASED ON IN VITRO DATA WITH AN EFFECTIVE AMOUNT OF ELX, TEZ, AND IVA
- U-3034 TREATMENT OF TRD IN ADULTS BY NASALLY ADMINISTERING 56MG OR 84MG OF ESKETAMINE 2X WEEKLY FOR 4 WEEKS DURING THE INDUCTION PHASE IN CONJUNCTION WITH AN ORAL ANTIDEPRESSANT
- U-3035 TREATMENT OF DEPRESSIVE SYMPTOMS IN ADULTS WITH MDD WITH ACUTE SUICIDAL IDEATION OR BEHAVIOR BY NASALLY ADMINISTERING 56MG OR 84 MG OF ESKETAMINE 2X WEEKLY FOR 4 WEEKS IN CONJUNCTION WITH AN ORAL ANTIDEPRESSANT
- U-3036 TREATMENT OF TRD IN ADULTS BY NASALLY ADMINISTERING 56MG OR 84MG OF ESKETAMINE 2X WEEKLY FOR 4 WEEKS DURING THE INDUCTION PHASE FOLLOWED BY A MAINTENANCE PHASE OF 56MG OR 84 MG WEEKLY OR 1X EVERY TWO WEEKS IN CONJUNCTION WITH AN ORAL ANTIDEPRESSANT
- U-3037 A METHOD OF DELIVERING NITRIC OXIDE TO A PATIENT
- U-3038 MANAGEMENT OF MODERATE-TO-SEVERE PAIN BY INTRAVENOUS INJECTION IN PATIENTS WITH MILD RENAL IMPAIRMENT
- U-3039 TREATMENT OF DUCHENNE MUSCULAR DYSTROPHY (DMD) IN PATIENTS WHO HAVE A CONFIRMED MUTATION OF THE DMD GENE THAT IS AMENABLE TO EXON 53 SKIPPING
- U-3040 LONG-TERM TREATMENT OF ADULTS WITH GAUCHER DISEASE TYPE 1 WHO ARE CYP2D6 EXTENSIVE METABOLIZERS WITH MILD HEPATIC IMPAIRMENT AND ARE CONCURRENTLY TAKING A STRONG OR MODERATE CYP3A INHIBITOR
- U-3041 LONG-TERM TREATMENT OF ADULTS WITH GAUCHER DISEASE TYPE 1 WHO ARE CYP2D6 EXTENSIVE METABOLIZERS WITH MODERATE TO SEVERE RENAL IMPAIRMENT
- U-3042 LONG-TERM TREATMENT OF ADULTS WITH GAUCHER DISEASE TYPE 1 WHO ARE CYP2D6 POOR METABOLIZERS WITH 84 MG ONCE DAILY OF ELIGLUSTAT (EQUIVALENT TO 100 MG OF ELIGLUSTAT TARTRATE)
- U-3043 LONG-TERM TREATMENT OF ADULTS WITH GAUCHER DISEASE TYPE 1 WHO ARE CYP2D6 EXTENSIVE OR INTERMEDIATE METABOLIZERS WITH 84 MG TWICE PER DAY OF ELIGLUSTAT

PATENT AND EXCLUSIVITY TERMS

ADB 181 of 225

PATENT USE

(EQUIVALENT TO 100 MG OF ELIGLUSTAT TARTRATE TWICE PER DAY)

- U-3044 AXITINIB IN COMBINATION WITH AVELUMAB FOR THE FIRST-LINE TREATMENT OF PATIENTS WITH ADVANCED RENAL CELL CARCINOMA
- U-3045 TREATMENT OF OVERACTIVE BLADDER (OAB) WITH SYMPTOMS OF URGE URINARY INCONTINENCE, URGENCY, AND URINARY FREQUENCY
- U-3046 METHOD OF ADMINISTERING VALBENAZINE WHILE AVOIDING CONCOMITANT USE OF A STRONG CYP3A4 INDUCER
- U-3047 USE IN COMBINATION WITH CAPECITABINE, FOR THE TREATMENT OF ADULT PATIENTS WITH ADVANCED OR METASTATIC HER2-POSITIVE BREAST CANCER WHO HAVE RECEIVED TWO OR MORE PRIOR ANTI-HER2 BASED REGIMENS IN THE METASTATIC SETTING
- U-3048 A METHOD OF TREATING IRON DEFICIENCY ANEMIA IN ADULT PATIENTS WHO HAVE INTOLERANCE TO ORAL IRON OR HAVE HAD UNSATISFACTORY RESPONSE TO ORAL IRON BY INTRAVENOUSLY ADMINISTERING FERRIC CARBOXYMALTOSE TO PROVIDE 500 TO 750 MG OF ELEMENTAL IRON
- U-3049 A METHOD OF TREATING IRON DEFICIENCY ANEMIA IN ADULT PATIENTS WHO HAVE NON-DIALYSIS DEPENDENT CHRONIC KIDNEY DISEASE BY INTRAVENOUSLY ADMINISTERING FERRIC CARBOXYMALTOSE TO PROVIDE 500 TO 750 MG OF ELEMENTAL IRON
- U-3050 METHOD OF TREATING IRON DEFICIENCY ANEMIA IN ADULT PATIENTS WHO HAVE INTOLERANCE TO ORAL IRON OR HAVE HAD UNSATISFACTORY RESPONSE TO ORAL IRON BY INTRAVENOUSLY ADMINISTERING AT LEAST ABOUT 0.6G OF IRON AS FERRIC CARBOXYMALTOSE IN ABOUT 15 MIN OR LESS
- U-3051 METHOD OF TREATING IRON DEFICIENCY ANEMIA IN ADULT PATIENTS WHO HAVE NON-DIALYSIS DEPENDENT CHRONIC KIDNEY DISEASE BY INTRAVENOUSLY ADMINISTERING AT LEAST ABOUT 0.6G OF IRON AS FERRIC CARBOXYMALTOSE IN ABOUT 15 MIN OR LESS
- U-3052 USE OF VASCEPA TO REDUCE TRIGLYCERIDE LEVELS IN AN ADULT PATIENT ON STATIN THERAPY AND HAVING ATRIAL FIBRILLATION AND TRIGLYCERIDE LEVELS OF GREATER THAN 500 MG/DL
- U-3053 USE OF VASCEPA AS AN ADJUNCT TO STATIN THERAPY TO REDUCE THE RISK OF MYOCARDIAL INFARCTION IN AN ADULT PATIENT WITH ELEVATED TRIGLYCERIDE LEVELS AND ESTABLISHED CV DISEASE OR DIABETES MELLITUS AND TWO OR MORE ADDITIONAL RISK FACTORS FOR CV DISEASE
- U-3054 TREATMENT OF DRUG-INDUCED EXTRAPYRAMIDAL REACTIONS IN ADULT PATIENTS WITH PARKINSON'S DISEASE
- U-3055 A METHOD OF TREATING HUNTINGTON'S CHOREA
- U-3056 TREATMENT OF PATIENTS WITH ACTIVE LUPUS NEPHRITIS
- U-3057 TREATMENT OF PATIENTS WITH METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) THAT IS ANAPLASTIC LYMPHOMA KINASE (ALK)-POSITIVE
- U-3058 TREATMENT OF PEDIATRIC PATIENTS 1 YEAR OF AGE AND OLDER AND YOUNG ADULTS WITH RELAPSED OR REFRACTORY, SYSTEMIC ANAPLASTIC LARGE CELL LYMPHOMA (ALCL) THAT IS ANAPLASTIC LYMPHOMA KINASE (ALK)-POSITIVE
- U-3059 TREATMENT OF HIV-1 INFECTION IN ADULTS TO REPLACE THE CURRENT ANTIRETROVIRAL REGIMEN IN THOSE WHO ARE VIROLOGICALLY SUPPRESSED ON A STABLE ANTIRETROVIRAL REGIMEN WITH NO HISTORY OF TREATMENT FAILURE
- U-3060 TREATMENT OF HIV INFECTION IN ADULTS
- U-3061 TREATMENT OF HIV-1 IN AN ADULT IN COMBINATION WITH RILPIVIRINE
- U-3062 REDUCING THE RISK OF CARDIOVASCULAR DEATH AND HEART FAILURE (HF) HOSPITALIZATION FOLLOWING A HOSPITALIZATION FOR HF OR NEED FOR OUTPATIENT IV DIURETICS, IN ADULTS WITH SYMPTOMATIC CHRONIC HF AND EJECTION FRACTION LESS THAN 45%
- U-3063 RELAPSED OR REFRACTORY MARGINAL ZONE LYMPHOMA (MZL) WHO HAVE RECEIVED AT LEAST ONE PRIOR ANTI-CD20-BASED REGIMEN
- U-3064 RELAPSED OR REFRACTORY FOLLICULAR LYMPHOMA (FL) WHO HAVE RECEIVED AT LEAST THREE PRIOR LINES OF SYSTEMIC THERAPY
- U-3065 TREATMENT OF ADULTS WITH METASTATIC UROTHELIAL CARCINOMA THAT HAS SUSCEPTIBLE FGFR3 OR FGFR2 GENETIC ALTERATIONS AND PROGRESSED DURING OR FOLLOWING PRIOR PLATINUM-CONTAINING CHEMOTHERAPY
- U-3066 TREATMENT OF ADULTS WITH LOCALLY ADVANCED OR METASTATIC, SURGICALLY UNRESECTABLE UROTHELIAL CARCINOMA THAT HAS SUSCEPTIBLE FGFR3 OR FGFR2 GENETIC ALTERATIONS AND PROGRESSED DURING OR FOLLOWING PRIOR PLATINUM-CONTAINING CHEMOTHERAPY

PATENT AND EXCLUSIVITY TERMS

ADB 182 of 225

PATENT USE

- U-3067 TREATMENT OF ADULTS WITH LOCALLY ADVANCED OR METASTATIC UROTHELIAL CARCINOMA THAT HAS SUSCEPTIBLE FGFR3 GENETIC ALTERATIONS AND PROGRESSED DURING OR FOLLOWING PRIOR PLATINUM-CONTAINING CHEMOTHERAPY
- U-3068 TREATMENT IN COMBINATION WITH CABOTEGRAVIR OF HIV-1 INFECTION IN ADULTS TO REPLACE THE CURRENT ANTIRETROVIRAL REGIMEN IN THOSE WHO ARE VIROLOGICALLY SUPPRESSED ON A STABLE ANTIRETROVIRAL REGIMEN WITH NO HISTORY OF TREATMENT FAILURE
- U-3069 TREATMENT OF ACUTE UNCOMPLICATED INFLUENZA IN PATIENTS 6 MONTHS AND OLDER
- U-3070 REDUCING THE RISK OF MORTALITY IN HUTCHINSON-GILFORD PROGERIA SYNDROME (HGPS)
- U-3071 USE FOR THE TREATMENT OF SEIZURES IN PATIENTS WITH TUBEROUS SCLEROSIS COMPLEX
- U-3072 USE FOR THE TREATMENT OF GENERALIZED SEIZURES OR FOCAL SEIZURES WITH IMPAIRMENT IN PATIENTS WITH TUBEROUS SCLEROSIS COMPLEX
- U-3073 USE FOR REDUCING SEIZURE FREQUENCY IN PATIENTS WITH TUBEROUS SCLEROSIS COMPLEX
- U-3074 METHOD FOR PROVIDING SUSTAINED LOCAL ANESTHESIA FOR AT LEAST 24 HOURS
- U-3075 TREATMENT OF ADRENAL INSUFFICIENCY
- U-3076 METHOD OF TREATING TARDIVE DYSKINESIA WHILE AVOIDING CONCOMITANT USE OF A STRONG CYP3A4 INDUCER
- U-3077 TREATING A SOLID TUMOR, INCLUDING LUNG CANCER, WITH A MET ALTERATION(S), OR STABILIZING OR IMPROVING SYMPTOMS ASSOCIATED WITH HAVING A SOLID TUMOR, INCLUDING LUNG CANCER, WITH A MET ALTERATION(S), BY ADMINISTERING AN EFFECTIVE AMOUNT OF TEPOTINIB
- U-3078 TREATING A SOLID TUMOR, INCLUDING LUNG CANCER, HAVING A MET KINASE ALTERATION(S) BY ADMINISTERING AN EFFECTIVE AMOUNT OF TEPOTINIB
- U-3079 A METHOD TO DECREASE THE INCIDENCE OF CHEMOTHERAPY-INDUCED MYELOSUPPRESSION IN ADULT PATIENTS WHEN ADMINISTERED PRIOR TO A PLATINUM/ETOPOSIDE-CONTAINING REGIMEN FOR EXTENSIVE-STAGE SMALL CELL LUNG CANCER
- U-3080 A METHOD TO DECREASE THE INCIDENCE OF CHEMOTHERAPY-INDUCED MYELOSUPPRESSION IN ADULT PATIENTS WHEN ADMINISTERED PRIOR TO A TOPOTECAN-CONTAINING REGIMEN FOR EXTENSIVE- STAGE SMALL CELL LUNG CANCER
- U-3081 A METHOD TO DECREASE THE INCIDENCE OF CHEMOTHERAPY-INDUCED MYELOSUPPRESSION IN ADULT PATIENTS WHEN ADMINISTERED PRIOR TO A CARBOPLATIN AND ETOPOSIDE-CONTAINING REGIMEN FOR EXTENSIVE-STAGE SMALL CELL LUNG CANCER
- U-3082 TREATMENT OF EXCESSIVE DAYTIME SLEEPINESS ASSOCIATED WITH OBSTRUCTIVE SLEEP APNEA (OSA) IN AN ADULT THROUGH A DOSING REGIMEN THAT INCLUDES ORAL ADMINISTRATION OF 75 MG ONCE DAILY FOR AT LEAST 3 DAYS FOLLOWED BY 150 MG ONCE DAILY
- U-3083 METHOD OF TREATING TRANSFUSIONAL IRON OVERLOAD
- U-3084 TREATMENT OF HEART FAILURE WITH PRESERVED EJECTION FRACTION
- U-3085 DOSE MODIFICATION FOR RENAL IMPAIRMENT
- U-3086 FOR HIGH-DOSE CONDITIONING TREATMENT PRIOR TO HEMATOPOIETIC PROGENITOR (STEM) CELL TRANSPLANTATION IN PATIENTS WITH MULTIPLE MYELOMA
- U-3087 TREATMENT OF DUCHENNE MUSCULAR DYSTROPHY (DMD) IN PATIENTS WHO HAVE A MUTATION OF THE DMD GENE THAT IS AMENABLE TO EXON 45 SKIPPING BY INDUCING EXON-SKIPPING OF EXON 45
- U-3088 TREATMENT OF DUCHENNE MUSCULAR DYSTROPHY (DMD) IN PATIENTS WHO HAVE A MUTATION OF THE DMD GENE THAT IS AMENABLE TO EXON 45 SKIPPING
- U-3089 TREATMENT OF DUCHENNE MUSCULAR DYSTROPHY (DMD) IN PATIENTS WHO HAVE A MUTATION OF THE DMD GENE THAT IS AMENABLE TO EXON 45 SKIPPING BY RESTORING AN MRNA READING FRAME TO INDUCE DYSTROPHIN PROTEIN PRODUCTION
- U-3090 TREATMENT OF HOSPITAL-ACQUIRED BACTERIAL PNEUMONIA IN PATIENTS WITH END-STAGE RENAL DISEASE ON HEMODIALYSIS
- U-3091 TREATMENT OF VENTILATOR-ASSOCIATED BACTERIAL PNEUMONIA IN PATIENTS WITH END-STAGE RENAL DISEASE ON HEMODIALYSIS
- U-3092 METHOD OF TREATING MOLYBDENUM COFACTOR DEFICIENCY TYPE A
- U-3093 IN COMBINATION WITH DEXAMETHASONE TO TREAT RELAPSED OR REFRACTORY MULTIPLE MYELOMA (REFRACTORY TO AT LEAST 1 PROTEASOME INHIBITOR, 1 IMMUNOMODULATORY AGENT, AND 1 ANTI-CD38 MAB) IN ADULTS WHO RECEIVED AT LEAST 4 PRIOR LINES OF THERAPY

PATENT AND EXCLUSIVITY TERMS

ADB 183 of 225

PATENT USE

- U-3094 TREATMENT OF ATTENTION-DEFICIT HYPERACTIVITY DISORDER (ADHD) WITH SERDEXMETHYLPHENIDATE AND DEXMETHYLPHENIDATE
- U-3095 TREATMENT OF HYPERLIPIDEMIA
- U-3096 TREATMENT OF ADULT PATIENTS WITH METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) WHOSE TUMORS ARE ANAPLASTIC LYMPHOMA KINASE (ALK)-POSITIVE AS DETECTED BY AN FDA-APPROVED TEST
- U-3097 EXTENDED ADJUVANT TREATMENT OF ADULT PATIENTS WITH EARLY-STAGE HUMAN EPIDERMAL GROWTH FACTOR RECEPTOR 2 (HER2)-POSITIVE BREAST CANCER, TO FOLLOW ADJUVANT TRASTUZUMAB BASED THERAPY
- U-3098 TREATMENT OF REFRACTORY EPILEPSY PATIENTS WITH FENFLURAMINE THAT REDUCES THE RISK OF CARDIOVASCULAR TOXICITY BY USING CARDIAC MONITORING AND RESTRICTED DISTRIBUTION
- U-3099 TREATMENT OF EXCESSIVE DAYTIME SLEEPINESS ASSOCIATED WITH OBSTRUCTIVE SLEEP APNEA (OSA) IN A PATIENT WITH MODERATE RENAL IMPAIRMENT
- U-3100 A METHOD OF TREATING ADULTS WITH RELAPSED OR REFRACTORY ADVANCED RENAL CELL CARCINOMA FOLLOWING TWO OR MORE PRIOR SYSTEMIC THERAPIES BY INHIBITING THE ANGIOGENESIS OF BLOOD VESSELS WITH A VASCULAR ENDOTHELIAL GROWTH FACTOR INHIBITOR
- U-3101 TREATMENT OF ACUTE BACTERIAL SKIN AND SKIN STRUCTURE INFECTIONS WITH A SINGLE DOSE OF 1200MG ORITAVANCIN OR ITS SINGLE DOSE EQUIVALENT
- U-3102 REDUCTION OF CIRCULATING LYMPHOCYTES IN TREATING RELAPSING FORMS OF MULTIPLE SCLEROSIS (MS), TO INCLUDE CLINICALLY ISOLATED SYNDROME, RELAPSING-REMITTING DISEASE, AND ACTIVE SECONDARY PROGRESSIVE DISEASE, IN ADULTS
- U-3103 TREATMENT OF RELAPSING FORMS OF MULTIPLE SCLEROSIS (MS), TO INCLUDE CLINICALLY ISOLATED SYNDROME, RELAPSING-REMITTING DISEASE, AND ACTIVE SECONDARY PROGRESSIVE DISEASE, IN ADULTS USING A DOSE TITRATION SCHEDULE FOLLOWED BY A MAINTENANCE DOSE
- U-3104 TREATMENT OF C. DIFFICILE-ASSOCIATED DIARRHEA
- U-3105 TREATMENT OF STAPHYLOCOCCAL ENTEROCOLITIS
- U-3106 TREATMENT OF NIGHTTIME SLEEP DISTURBANCES IN SMITH-MAGENIS SYNDROME BY AVOIDING THE ADMINISTRATION OF TASIMELTEON TO SMOKERS OR TO PATIENTS BEING TREATED WITH A CYP1A2 INHIBITOR
- U-3107 TREATMENT OF NON-24 HOUR SLEEP-WAKE DISORDER BY AVOIDING THE ADMINISTRATION OF TASIMELTEON TO SMOKERS OR TO PATIENTS BEING TREATED WITH A CYP1A2 INHIBITOR
- U-3108 TREATMENT OF NEUROGENIC DETRUSOR OVERACTIVITY (NDO) IN PEDIATRIC PATIENTS AGED 3 YEARS AND OLDER BY ADMINISTRATION OF AN EXTENDED-RELEASE SUSPENSION FORMULATION OF MIRABEGRON
- U-3109 METHOD OF USING VISMODEGIB TO TREAT BASAL CELL CARCINOMA
- U-3110 USE OF NALOXONE FOR THE EMERGENCY TREATMENT OF KNOWN OR SUSPECTED OPIOID OVERDOSE, AS MANIFESTED BY RESPIRATORY AND/OR CENTRAL NERVOUS SYSTEM DEPRESSION, FOR ADULT AND PEDIATRIC PATIENTS
- U-3111 TREATING OPIOID USE DISORDER
- U-3112 TREATING NEWLY DIAGNOSED ACUTE MYELOGENOUS LEUKEMIA (AML) CHARACTERIZED BY THE PRESENCE OF A MUTANT ALLELE OF IDH1 BY ADMINISTERING A ONCE DAILY 500 MG ORAL DOSE TO A SUBJECT THAT HAS NOT INGESTED A HIGH-FAT MEAL
- U-3113 TREATING RELAPSED/REFRACTORY ACUTE MYELOGENOUS LEUKEMIA (AML) CHARACTERIZED BY THE PRESENCE OF A MUTANT ALLELE OF IDH1 BY ADMINISTERING A ONCE DAILY 500 MG ORAL DOSE TO A SUBJECT THAT HAS NOT INGESTED A HIGH-FAT MEAL
- U-3114 TREATMENT OF CHRONIC LYMPHOCYTIC LEUKEMIA (CLL) OR SMALL LYMPHOCYTIC LYMPHOMA (SLL) BY ORALLY ADMINISTERING VENETOCLAX TO AN ADULT ACCORDING TO A DOSE RAMP-UP THAT INCLUDES A DOSE OF 50 MG PER DAY FOR 1 WEEK FOLLOWED BY 100 MG PER DAY FOR 1 WEEK
- U-3115 METHOD OF TREATING IRON DEFICIENCY ANEMIA IN ADULT PATIENTS WHO HAVE INTOLERANCE TO ORAL IRON OR HAVE HAD UNSATISFACTORY RESPONSE TO ORAL IRON BY INTRAVENOUSLY ADMINISTERING ABOUT 1 G OF IRON AS FERRIC CARBOXYMALTOSE IN ABOUT 15 MINUTES OR LESS
- U-3116 METHOD OF TREATING IRON DEFICIENCY ANEMIA IN ADULT PATIENTS WHO HAVE NON-DIALYSIS DEPENDENT CHRONIC KIDNEY DISEASE BY INTRAVENOUSLY ADMINISTERING ABOUT 1 G OF IRON AS FERRIC CARBOXYMALTOSE IN ABOUT 15 MINUTES OR LESS

PATENT AND EXCLUSIVITY TERMS

ADB 184 of 225

PATENT USE

U-3117 ADMINISTRATION TO THE EYE OF A PATIENT FOR TREATMENT OF DRY EYE CONDITION

U-3118 TREATMENT OF POSTSURGICAL PAIN PROVIDING ANALGESIA TO A PATIENT FOR UP TO 72 HOURS, FOR EXAMPLE, AFTER BUNIONECTOMY, OPEN INGUINAL HERNIORRHAPHY, OR TOTAL KNEE ARTHROPLASTY VIA SOFT TISSUE OR PERIARTICULAR INSTILLATION

U-3119 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETIS BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 1 AND 11

U-3120 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETIS BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 1, 11, AND 12

U-3121 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETIS BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 13 AND 23

U-3122 ADJUNCT TO DIET EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETIS BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 13, 23, AND 24

U-3123 TREATMENT OF ADULT PATIENTS WITH PAROXYSMAL NOCTURNAL HEMOGLOBINURIA (PNH) BY ADMINISTRATION OF COMPLEMENT INHIBITOR PEGCETACOPLAN

U-3124 TREATMENT OF ADULT PATIENTS WITH PAROXYSMAL NOCTURNAL HEMOGLOBINURIA (PNH) BY ADMINISTRATION OF PEGCETACOPLAN

U-3125 USE FOR LOADING DOSE IN PATIENTS WITH SYMPTOMATIC AFIB/AFL WHO ARE CURRENTLY IN SINUS RHYTHM OR FOR THE TREATMENT OF LIFE-THREATENING VENTRICULAR TACHYCARDIA

U-3126 USE OF VASCEPA AS AN ADJUNCT TO STATIN THERAPY TO REDUCE THE RISK OF A SECOND AND FURTHER CARDIOVASCULAR EVENT IN AN ADULT PATIENT WITH ESTABLISHED CARDIOVASCULAR DISEASE

U-3127 REDUCTION OF THE RISK OF CARDIOVASCULAR DEATH AND HOSPITALIZATION FOR HEART FAILURE IN ADULTS WITH HEART FAILURE WITH REDUCED EJECTION FRACTION AND WITHOUT TYPE II DIABETES

U-3128 TREATMENT OF COMPLICATED URINARY TRACT INFECTIONS (CUTI) INCLUDING PYELONEPHRITIS CAUSED BY THE FOLLOWING SUSCEPTIBLE MICROORGANISMS: ESCHERICHIA COLI, KLEBSIELLA PNEUMONIA, ENTEROBACTER CLOACAE SPECIES COMPLEX WITH MEROPENEM & VABORBACTAM AS SPECIFIED

U-3129 MANAGEMENT OF HEAVY MENSTRUAL BLEEDING ASSOCIATED WITH UTERINE LEIOMYOMAS (FIBROIDS) IN PREMENOPAUSAL WOMEN

U-3130 METHOD OF POSITRON EMISSION TOMOGRAPHY (PET) IN MEN WITH PROSTATE CANCER

U-3131 USE OF ZUBSOLV FOR TREATMENT OF OPIOID DEPENDENCE

U-3132 INDICATED FOR THE TREATMENT OF MODERATELY TO SEVERELY ACTIVE ULCERATIVE COLITIS

U-3133 TREATING SICKLE CELL DISEASE BY ADMINISTERING 1500 MG OF VOXELOTOR ORALLY ONCE DAILY

U-3134 INCREASING HEMOGLOBIN TO TREAT SICKLE CELL DISEASE BY ADMINISTERING 1500 MG OF VOXELOTOR ORALLY ONCE DAILY

U-3135 TREATING SCHIZOPHRENIA

U-3136 METHOD OF TREATING SCHIZOPHRENIA BY ADMINISTERING THE ATYPICAL ANTIPSYCHOTIC OLANZAPINE AND SAMIDORPHAN, WITH REDUCED ANTIPSYCHOTIC INDUCED WEIGHT GAIN

U-3137 METHOD OF TREATING BIPOLAR DISORDER BY ADMINISTERING THE ATYPICAL ANTIPSYCHOTIC OLANZAPINE AND SAMIDORPHAN, WITH REDUCED ANTIPSYCHOTIC INDUCED WEIGHT GAIN

U-3138 METHOD OF TREATING SCHIZOPHRENIA BY ADMINISTERING THE ATYPICAL ANTIPSYCHOTIC OLANZAPINE AND SAMIDORPHAN, WITH REDUCTION OF THE ADVERSE METABOLIC PROFILE

U-3139 METHOD OF TREATING BIPOLAR DISORDER BY ADMINISTERING THE ATYPICAL ANTIPSYCHOTIC OLANZAPINE AND SAMIDORPHAN, WITH REDUCTION OF THE ADVERSE METABOLIC PROFILE

U-3140 METHOD OF TREATING SCHIZOPHRENIA BY ADMINISTERING THE ATYPICAL ANTIPSYCHOTIC OLANZAPINE AND SAMIDORPHAN

U-3141 METHOD OF TREATING BIPOLAR DISORDER BY ADMINISTERING THE ATYPICAL ANTIPSYCHOTIC OLANZAPINE AND SAMIDORPHAN

U-3142 PREVENTIVE TREATMENT OF EPISODIC MIGRAINE IN ADULTS

U-3143 FOR THE TOPICAL TREATMENT OF PLAQUE PSORIASIS IN PATIENTS 18 YEARS OF AGE AND OLDER

PATENT AND EXCLUSIVITY TERMS

ADB 185 of 225

PATENT USE

- U-3144 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 6 YEARS AND OLDER WHO HAVE AT LEAST ONE F508DEL MUTATION IN THE CFTR GENE OR A MUTATION IN THE CFTR GENE THAT IS RESPONSIVE BASED ON IN VITRO DATA WITH ELEXACAFTOR, TEZACAFTOR, AND IVACAFTOR
- U-3145 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 6 YEARS AND OLDER WHO HAVE A R117H MUTATION IN THE CFTR GENE WITH ELEXACAFTOR, TEZACAFTOR, AND IVACAFTOR
- U-3146 TREATMENT OF A MODERATE TO MILD CLINICAL PHENOTYPE OF CF IN PATIENTS AGED 6 YEARS AND OLDER WHO HAVE AT LEAST ONE F508DEL MUTATION IN THE CFTR GENE OR A MUTATION IN THE CFTR GENE THAT IS RESPONSIVE BASED ON IN VITRO DATA WITH ELX, TEZ, AND IVA
- U-3147 FOR THE TREATMENT OF NEWLY-DIAGNOSED THERAPY-RELATED ACUTE MYELOID LEUKEMIA (T-AML) OR AML WITH MYELODYSPLASIA-RELATED CHANGES (AML-MRC) IN ADULTS AND PEDIATRIC PATIENTS 1 YEAR AND OLDER
- U-3148 METHOD OF TREATING RELAPSING FORMS OF MULTIPLE SCLEROSIS BEFORE AND AFTER ADMINISTERING AN INACTIVE VACCINE
- U-3149 METHOD OF RECONSTITUTING A LYOPHILIZED LIPOSOMAL COMPOSITION FOR ADMINISTERING CYTARABINE AND DAUNORUBICIN TO TREAT NEWLY-DIAGNOSED THERAPY-RELATED AML (T-AML) OR AML WITH MYELODYSPLASIA-RELATED CHANGES (AML-MRC) IN PATIENTS 1 YEAR AND OLDER
- U-3150 METHOD OF ADMINISTERING A RECONSTITUTED LIPOSOMAL COMPOSITION CONTAINING CYTARABINE AND DAUNORUBICIN TO TREAT NEWLY-DIAGNOSED THERAPY-RELATED AML (T-AML) OR AML WITH MYELODYSPLASIA-RELATED CHANGES (AML-MRC) IN PATIENTS 1 YEAR AND OLDER
- U-3151 TO IMPROVE WAKEFULNESS IN ADULT PATIENTS WITH EXCESSIVE DAYTIME SLEEPINESS ASSOCIATED WITH NARCOLEPSY OR OBSTRUCTIVE SLEEP APNEA (OSA) WITH A DOSING REGIMEN THAT INCLUDES A DOSE OF 75 MG ONCE DAILY FOR AT LEAST 3 DAYS FOLLOWED BY 150 MG ONCE DAILY
- U-3152 USE BY FEMALES OF REPRODUCTIVE POTENTIAL TO PREVENT PREGNANCY
- U-3153 TREATMENT OF ADVANCED GASTROINTESTINAL STROMAL TUMOR IN PATIENTS HAVING PROGRESSED FROM A FIRST LINE ADMINISTRATION OF IMATINIB, A SECOND LINE ADMINISTRATION OF SUNITINIB, AND A THIRD LINE ADMINISTRATION OF REGORAFENIB
- U-3154 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 6 YEARS AND OLDER WHO HAVE AT LEAST ONE F508DEL MUTATION IN THE CFTR GENE WITH A COMPOSITION COMPRISING ELEXACAFTOR, TEZACAFTOR, AND IVACAFTOR; AND ANOTHER COMPOSITION COMPRISING IVACAFTOR
- U-3155 TREATMENT OF CF IN PATIENTS 6 YEARS AND OLDER WHO HAVE AT LEAST ONE F508DEL MUTATION OR A MUTATION IN THE CFTR GENE THAT IS RESPONSIVE BASED ON IN VITRO DATA USING A SOLID COMPOSITION COMPRISING ELX, TEZ, AMORPHOUS IVA, AND < ~30% CRYSTALLINE IVA
- U-3156 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 6 YEARS AND OLDER WHO HAVE AT LEAST ONE F508DEL MUTATION IN THE CFTR GENE WITH AN EFFECTIVE AMOUNT OF A PHARMACEUTICAL COMPOSITION COMPRISING ELEXACAFTOR, TEZACAFTOR, AND IVACAFTOR
- U-3157 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 6 YEARS AND OLDER WHO HAVE AT LEAST ONE F508DEL MUTATION IN THE CFTR GENE OR A MUTATION IN THE CFTR GENE THAT IS RESPONSIVE BASED ON IN VITRO DATA WITH A COMPOSITION ACCORDING TO CLAIM 1 OF US 10081621
- U-3158 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 6 YEARS AND OLDER WHO HAVE AT LEAST ONE F508DEL MUTATION IN THE CFTR GENE OR A MUTATION IN THE CFTR GENE THAT IS RESPONSIVE BASED ON IN VITRO DATA WITH AN EFFECTIVE AMOUNT OF ELX, TEZ, AND IVA
- U-3159 TREATMENT OF ADULT AND POST-MENARCHAL PEDIATRIC FEMALES WITH VULVOVAGINAL CANDIDIASIS (VVC)
- U-3160 PROPHYLAXIS OF INVASIVE ASPERGILLUS AND CANDIDA INFECTIONS IN ADULTS AND PEDIATRIC PATIENTS 2 YEARS OF AGE AND OLDER WHO ARE SEVERELY IMMUNOCOMPROMISED
- U-3161 METHOD FOR WEIGHT MANAGEMENT ACCORDING TO A DOSE ESCALATION SCHEDULE
- U-3162 METHOD FOR WEIGHT MANAGEMENT
- U-3163 TREATMENT OF ACTINIC KERATOSES OF UPPER EXTREMITIES BY PHOTODYNAMIC THERAPY
- U-3164 GASTROINTESTINAL TABLETS INDICATED FOR CLEANSING THE COLON IN PREPARATION FOR COLONOSCOPY
- U-3165 METHOD OF TREATING HUMAN SMALLPOX DISEASE

PATENT AND EXCLUSIVITY TERMS

ADB 186 of 225

PATENT USE

- U-3166 OTC USE: ALLERGY SYMPTOM RELIEVER; TEMPORARY RELIEF OF THESE SYMPTOMS DUE TO HAY FEVER OR OTHER UPPER RESPIRATORY ALLERGIES: NASAL CONGESTION, RUNNY NOSE, SNEEZING AND ITCHY NOSE
- U-3167 TREATMENT OF CHRONIC HEPATITIS C VIRUS (HCV) GENOTYPE 1, 2, 3, 4, OR 6 IN PEDIATRIC PATIENTS 3 TO LESS THAN 12 YEARS OF AGE OR WEIGHING LESS THAN 45 KG
- U-3168 TREATMENT OF ADVANCED SYSTEMIC MASTOCYTOSIS, INCLUDING PATIENTS WITH AGGRESSIVE SYSTEMIC MASTOCYTOSIS (ASM), SYSTEMIC MASTOCYTOSIS WITH AN ASSOCIATED HEMATOLOGICAL NEOPLASM (SM-AHN), AND MAST CELL LEUKEMIA (MCL)
- U-3169 TREATMENT OF TRICHOMONIASIS IN ADULTS
- U-3170 TREATING CHRONIC HEART FAILURE WITH REDUCED EJECTION FRACTION IN PATIENTS NOT TAKING AN ACE INHIBITOR OR AN ARB OR PREVIOUSLY TAKING LOW DOSES OF THESE AGENTS, BY TITRATING UP FROM HALF THE USUALLY RECOMMENDED STARTING DOSE
- U-3171 TREATMENT OF INVASIVE ASPERGILLOSIS IN ADULTS AND PEDIATRIC PATIENTS 13 YEARS OF AGE AND OLDER
- U-3172 TREATMENT OF ADULT PATIENTS WITH PAROXYSMAL NOCTURNAL HEMOGLOBINURIA (PNH) BY ADMINISTRATION OF 1080 MG OF PEGCETACOPLAN
- U-3173 TREATMENT OF ADULT PATIENTS WITH PAROXYSMAL NOCTURNAL HEMOGLOBINURIA (PNH) BY ADMINISTRATION OF 1080 MG OF PEGCETACOPLAN TWICE WEEKLY
- U-3174 TREATMENT OF ADULT PATIENTS WITH PAROXYSMAL NOCTURNAL HEMOGLOBINURIA (PNH) BY ADMINISTRATION OF 1080 MG OF PEGCETACOPLAN EVERY THREE DAYS
- U-3175 TREATMENT OF COMPLICATED SKIN AND SKIN STRUCTURE INFECTIONS AND STAPHYLOCOCCUS AUREUS BLOODSTREAM INFECTIONS (BACTEREMIA) INCLUDING THOSE WITH RIGHT-SIDED INFECTIVE ENDOCARDITIS BY ADMINISTERING THE FORMULATION OF DAPTOMYCIN AS RECITED IN CLAIM 18
- U-3176 TREATMENT OF COMPLICATED SKIN AND SKIN STRUCTURE INFECTIONS AND S. AUREUS BLOODSTREAM INFECTIONS (BACTEREMIA) INCLUDING THOSE WITH RIGHT-SIDED INFECTIVE ENDOCARDITIS BY RECONSTITUTING AND ADMINISTERING THE FORMULATION AS RECITED IN CLAIM 12
- U-3177 TREATMENT OF VENOUS THROMBOTIC DISEASE
- U-3178 REDUCE THE RISK OF RECURRENCE OF VENOUS THROMBOTIC DISEASE
- U-3179 METHOD OF TREATING FOLLICULAR LYMPHOMA
- U-3180 DECREASING OFF TIME IN PATIENTS WITH PARKINSON'S DISEASE RECEIVING LEVODOPA-BASED THERAPY, WITH OR WITHOUT CONCOMITANT DOPAMINERGIC MEDICATIONS
- U-3181 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS 6 YEARS AND OLDER WHO ARE HOMOZYGOUS FOR THE F508DEL MUTATION IN THE CFTR GENE USING THE TABLET ACCORDING TO CLAIM 1 OF U.S. PATENT NO. 11,052,075, WHERE THE TABLET FURTHER COMPRISES IVACAFTOR
- U-3182 METHOD OF PROVIDING POSTSURGICAL PAIN MANAGEMENT, FOR EXAMPLE, VIA INFILTRATION FOR LOCAL ANALGESIA OR VIA INTERSCALENE BRACHIAL PLEXUS NERVE BLOCK FOR REGIONAL ANALGESIA
- U-3183 USE OF EPHEDRINE SULFATE FOR TREATING HYPOTENSION
- U-3184 MEKINIST(R) IS INDICATED, IN COMBINATION WITH DABRAFENIB, FOR THE ADJUVANT TREATMENT OF PATIENTS WITH MELANOMA WITH BRAF V600E OR V600K MUTATIONS AS DETECTED BY AN FDA-APPROVED TEST, AND INVOLVEMENT OF LYMPH NODE(S), FOLLOWING COMPLETE RESECTION
- U-3185 TAFINLAR(R) IS INDICATED, IN COMBINATION WITH TRAMETINIB, FOR THE ADJUVANT TREATMENT OF PATIENTS WITH MELANOMA WITH BRAF V600E OR V600K MUTATIONS AS DETECTED BY AN FDA-APPROVED TEST, AND INVOLVEMENT OF LYMPH NODE(S), FOLLOWING COMPLETE RESECTION
- U-3186 METHOD OF TREATING PRURITUS IN PATIENTS 3 MONTHS OR OLDER SUFFERING FROM PROGRESSIVE FAMILIAL INTRAHEPATIC CHOLESTASIS (PFIC)
- U-3187 METHOD OF REDUCING SERUM BILE ACIDS IN PATIENTS 3 MONTHS OR OLDER SUFFERING FROM PROGRESSIVE FAMILIAL INTRAHEPATIC CHOLESTASIS (PFIC)
- U-3188 IMPROVING GLYCEMIC CONTROL IN PATIENTS 10 YEARS OF AGE AND OLDER WITH TYPE 2 DIABETES MELLITUS BY ADMINISTERING A SUSTAINED-RELEASE EXENATIDE FORMULATION AS AN ADJUNCT TO DIET AND EXERCISE
- U-3189 AS AN ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN PATIENTS 10 TO 17 YEARS OF AGE WITH TYPE 2 DIABETES MELLITUS IN COMBINATION WITH INSULIN ALONE OR INSULIN PLUS ONE OTHER ORAL ANTIDIABETIC MEDICATION

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

- U-3190 AS AN ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN PATIENTS 10 TO 17 YEARS OF AGE WITH TYPE 2 DIABETES MELLITUS IN COMBINATION WITH METFORMIN AND/OR SULFONYLUREA
- U-3191 METHOD OF TREATING TYPE 2 DIABETES MELLITUS IN A PATIENT WITH RENAL IMPAIRMENT (30 ML/MIN/1.73 M² ≤ EGFR < 60 ML/MIN/1.73 M²) BY ONCE DAILY ADMINISTRATION OF 10 MG OR 25 MG OF EMPAGLIFLOZIN
- U-3192 METHOD OF TREATING TYPE 2 DIABETES MELLITUS IN A PATIENT WITH RENAL IMPAIRMENT (EGFR < 60 ML/MIN/1.73 M²) BY INITIATION OF EMPAGLIFLOZIN, LINAGLIPTIN AND METFORMIN HCL IF EGFR ≥ 45 ML/MIN/1.73 M² AND DISCONTINUATION IF EGFR < 30 ML/MIN/1.73 M²
- U-3193 METHOD OF TREATING TYPE 2 DIABETES MELLITUS IN A PATIENT WITH RENAL IMPAIRMENT (EGFR < 60 ML/MIN/1.73 M²) BY INITIATION OF EMPAGLIFLOZIN AND METFORMIN HCL IF EGFR ≥ 45 ML/MIN/1.73 M² AND DISCONTINUATION IF EGFR < 30 ML/MIN/1.73 M²
- U-3194 TOPICAL TREATMENT OF ACNE VULGARIS IN ADULTS AND PEDIATRIC PATIENTS 9 YEARS OF AGE AND OLDER
- U-3195 TREATMENT OF ADULTS WITH REFRACTORY, MODERATE-TO-SEVERE ATOPIC DERMATITIS NOT ADEQUATELY CONTROLLED WITH OTHER SYSTEMIC DRUG PRODUCTS OR WHEN USE OF THOSE THERAPIES IS INADVISABLE
- U-3196 TREATMENT OF ADULTS WITH LOCALLY ADVANCED OR METASTATIC UROTHELIAL CARCINOMA WITH SUSCEPTIBLE FGFR3 OR FGFR2 GENETIC ALTERATIONS, AND PROGRESSED DURING OR FOLLOWING PRIOR PLATINUM-CONTAINING CHEMOTHERAPY, WITH DOSING BASED ON SERUM PHOSPHATE LEVELS
- U-3197 MANAGEMENT OF PAIN SEVERE ENOUGH TO REQUIRE AN OPIOID ANALGESIC AND FOR WHICH ALTERNATIVE TREATMENTS ARE INADEQUATE
- U-3198 METHOD OF TREATING PATIENTS WITH IDIOPATHIC HYPERSOMNIA WITH SODIUM OXYBATE WHEN DIVALPROEX SODIUM IS CONCOMITANTLY ADMINISTERED
- U-3199 METHOD FOR REDUCING THE RISK OF CARDIOVASCULAR DEATH PLUS HOSPITALIZATION FOR HEART FAILURE IN ADULTS WITH TYPE 2 DIABETES MELLITUS, HEART FAILURE AND REDUCED EJECTION FRACTION BY ONCE DAILY ADMINISTRATION OF EMPAGLIFLOZIN
- U-3200 INCREASING SURVIVAL IN MCRPC PATIENTS PREVIOUSLY TREATED WITH DOCETAXEL BY ADMINISTERING CABAZITAXEL IN COMBINATION WITH PREDNISONE OR PREDNISOLONE AFTER A PREMEDICATION REGIMEN THAT INCLUDES AN ANTIHISTAMINE, A CORTICOSTEROID, AND AN H₂-ANTAGONIST
- U-3201 TREATMENT OF ADULT PATIENTS WITH VON HIPPEL-LINDAU DISEASE WHO REQUIRE THERAPY FOR ASSOCIATED RENAL CELL CARCINOMA, CENTRAL NERVOUS SYSTEM HEMANGIOBLASTOMAS, OR PANCREATIC NEUROENDOCRINE TUMORS, NOT REQUIRING IMMEDIATE SURGERY
- U-3202 MAINTENANCE TREATMENT OF CHRONIC OBSTRUCTIVE PULMONARY DISEASE (COPD) COMPRISING THE ONCE PER DAY ADMINISTRATION OF TRELEGY ELLIPTA, 100 MCG FLUTICASONE FUROATE/62.5 MCG UMECLIDINIUM/25 MCG VILANTEROL
- U-3203 MAINTENANCE TREATMENT OF PATIENTS WITH CHRONIC OBSTRUCTIVE PULMONARY DISEASE (COPD)
- U-3204 TREATMENT OF MODERATE-TO-SEVERE PRURITUS ASSOCIATED WITH CHRONIC KIDNEY DISEASE (CKD-AP) IN ADULTS UNDERGOING HEMODIALYSIS (HD)
- U-3205 REDUCTION OF RISK OF MAJOR CARDIOVASCULAR EVENTS (CARDIOVASCULAR DEATH, MYOCARDIAL INFARCTION AND STROKE) IN PATIENTS WITH CAD
- U-3206 REDUCTION OF RISK OF MAJOR THROMBOTIC VASCULAR EVENTS (MYOCARDIAL INFARCTION, ISCHEMIC STROKE, ACUTE LIMB ISCHEMIA, AND MAJOR AMPUTATION OF VASCULAR ETIOLOGY) IN PATIENTS WITH PAD
- U-3207 REDUCTION OF RISK OF CARDIOVASCULAR DEATH, MYOCARDIAL INFARCTION, AND STROKE IN PATIENTS WITH CAD BY ADMINISTERING CLINICALLY PROVEN EFFECTIVE AMOUNTS THAT ARE 2.5 MG RIVAROXABAN TWICE DAILY AND 75-100 MG ASPIRIN DAILY
- U-3208 REDUCTION OF RISK OF MYOCARDIAL INFARCTION AND ISCHEMIC STROKE IN PATIENTS WITH PAD BY ADMINISTERING CLINICALLY PROVEN EFFECTIVE AMOUNTS THAT ARE 2.5 MG RIVAROXABAN TWICE DAILY AND 75-100 MG ASPIRIN DAILY
- U-3209 USE OF VASCEPA AS AN ADJUNCT TO STATIN THERAPY TO REDUCE THE RISK MYOCARDIAL INFARCTION IN AN ADULT PATIENT HAVING ATRIAL FIBRILLATION OR ATRIAL FLUTTER AND ELEVATED TRIGLYCERIDE LEVELS
- U-3210 ONCE DAILY TREATMENT OF ANXIETY DISORDER IN ADULTS
- U-3211 TREATING DISTRIBUTIVE SHOCK WITH ANGIOTENSIN II

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-3212 TREATING SEPTIC SHOCK WITH ANGIOTENSIN II

U-3213 A METHOD OF TREATING A CANCER CHARACTERIZED BY AN IDH1 MUTATION WHEREIN THE CANCER IS PREVIOUSLY TREATED, LOCALLY ADVANCED OR METASTATIC CHOLANGIOCARCINOMA

U-3214 A METHOD OF TREATING PREVIOUSLY TREATED, LOCALLY ADVANCED OR METASTATIC CHOLANGIOCARCINOMA CHARACTERIZED BY THE PRESENCE OF A MUTANT ALLELE OF IDH1 BY ADMINISTERING A ONCE DAILY 500 MG ORAL DOSE TO A SUBJECT THAT HAS NOT INGESTED A HIGH-FAT MEAL

U-3215 A METHOD OF TREATING PREVIOUSLY TREATED, LOCALLY ADVANCED OR METASTATIC CHOLANGIOCARCINOMA CHARACTERIZED BY AN IDH1 MUTATION

U-3216 A METHOD FOR TREATING A BCRABL POSITIVE LEUKEMIA IN A SUBJECT THAT IS RESISTANT TO IMATINIB COMPRISING ADMINISTERING TO THE SUBJECT A THERAPEUTICALLY EFFECTIVE AMOUNT OF BOSUTINIB, WHEREIN THE SUBJECT HAS A MUTATION IN THE BCRABL PROTEIN AT 949T>C

U-3217 A METHOD FOR TREATING A BCRABL POSITIVE LEUKEMIA IN A SUBJECT THAT IS RESISTANT TO IMATINIB COMPRISING ADMINISTERING TO THE SUBJECT A THERAPEUTICALLY EFFECTIVE AMOUNT OF BOSUTINIB, WHEREIN THE SUBJECT HAS A MUTATION IN THE BCRABL PROTEIN AT F317L

U-3218 NASAL ADMINISTRATION OF DIHYDROERGOTAMINE MESYLATE BY METERED SPRAY FOR THE ACUTE TREATMENT OF MIGRAINE WITH OR WITHOUT AURA

U-3219 TREATMENT OF GASTROINTESTINAL STROMAL TUMOR

U-3220 TREATMENT OF PATIENTS WITH NON-SMALL CELL LUNG CANCER (NSCLC) WITH EPIDERMAL GROWTH FACTOR RECEPTOR (EGFR) EXON 20 EXON INSERTION MUTATIONS WHOSE DISEASE HAS PROGRESSED ON OR AFTER PLATINUM-BASED CHEMOTHERAPY

U-3221 USE OF VASCEPA AS AN ADJUNCT TO STATIN THERAPY TO REDUCE THE RISK OF A CARDIOVASCULAR EVENT IN A PATIENT WITH PRIOR PERCUTANEOUS CORONARY INTERVENTION

U-3222 TREATMENT OF ACUTE MYELOID LEUKEMIA (AML) BY ORALLY ADMINISTERING VENETOCLAX WITH AZACITIDINE OR DECITABINE OR LOW-DOSE CYTARABINE IN ADULTS 75 YEARS OR OLDER OR HAVING CERTAIN COMORBIDITIES ACCORDING TO A DOSE RAMP-UP INCLUDING A 100 MG PER DAY DOSE

U-3223 TREATMENT OF CHRONIC LYMPHOCYTIC LEUKEMIA (CLL) OR SMALL LYMPHOCYTIC LYMPHOMA (SLL) BY ORALLY ADMINISTERING VENETOCLAX TO AN ADULT ACCORDING TO A DOSE RAMP-UP INCLUDING A 100 MG PER DAY DOSE

U-3224 A METHOD OF TREATING HETEROZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA OR ESTABLISHED ATHEROSCLEROTIC CARDIOVASCULAR DISEASE BY DECREASING THE LEVEL OF LDL-C USING A FIXED DOSE COMBINATION OF 180 MG BEMPEDOIC ACID AND 10 MG EZETIMIBE

U-3225 TREATMENT OF DIFFERENTIATED THYROID CANCER THAT HAS PROGRESSED FOLLOWING PRIOR VEGFR-TARGETED THERAPY

U-3226 FOR TREATMENT OF CHRONIC GRAFT-VERSUS-HOST DISEASE (CGVHD) AFTER FAILURE OF ONE OR TWO LINES OF SYSTEMIC THERAPY

U-3227 FOR TREATMENT OF INTERMEDIATE OR HIGH-RISK MYELOFIBROSIS (MF), INCLUDING PRIMARY MF, POST-POLYCYTHEMIA VERA MF AND POST-ESSENTIAL THROMBOCYTHEMIA MF

U-3228 FOR TREATMENT OF POLYCYTHEMIA VERA (PV) IN PATIENTS WHO HAVE HAD AN INADEQUATE RESPONSE TO OR ARE INTOLERANT OF HYDROXYUREA

U-3229 FOR TOPICAL SHORT-TERM, NON-CONTINUOUS CHRONIC TREATMENT OF MILD TO MODERATE ATOPIC DERMATITIS IN NON-IMMUNOCOMPROMISED PATIENTS WHOSE DISEASE IS NOT ADEQUATELY CONTROLLED WITH TOPICAL PRESCRIPTION THERAPIES OR WHEN THOSE THERAPIES ARE NOT ADVISABLE

U-3230 FOR TREATMENT OF STEROID-REFRACTORY ACUTE GRAFT-VERSUS-HOST DISEASE (AGVHD)

U-3231 TREATMENT OF PHILADELPHIA CHROMOSOME POSITIVE CHRONIC MYELOID LEUKEMIA (PH+CML) BY ADMINISTERING NILOTINIB DISPERSED IN A FRUIT PREPARATION

U-3232 USE OF ORAL OCTREOTIDE FOR LONG-TERM MAINTENANCE TREATMENT IN ACROMEGALY PATIENTS WHILE AVOIDING CONCOMITANT ADMINISTRATION OF LEVONORGESTREL

U-3233 USE FOR THE TREATMENT OF SEIZURES ASSOCIATED WITH TUBEROUS SCLEROSIS COMPLEX IN PATIENTS TAKING EVEROLIMUS

U-3234 TREATMENT OF MACULAR EDEMA ASSOCIATED WITH UVEITIS

U-3235 USE FOR THE TREATMENT OF SEIZURES ASSOCIATED WITH DRAVET SYNDROME

U-3236 USE FOR THE TREATMENT OF SEIZURES ASSOCIATED WITH LENNOX-GASTAUT SYNDROME

U-3237 TREATMENT OF CHRONIC HEPATITIS C VIRUS (HCV) GENOTYPE 1, 2, 3, 4, 5, OR 6 IN

PATENT AND EXCLUSIVITY TERMS

ADB 189 of 225

PATENT USE

- ADULT AND PEDIATRIC PATIENTS 12 YEARS AND OLDER OR WEIGHING AT LEAST 45 KG
- U-3238 TREATMENT OF CHRONIC HEPATITIS C VIRUS (HCV) GENOTYPE 1, 2, 3, 4, 5, OR 6 IN PEDIATRIC PATIENTS 3 TO LESS THAN 12 YEARS OF AGE OR WEIGHING LESS THAN 45 KG
- U-3239 TREATMENT OF ADVANCED RENAL CELL CARCINOMA (RCC) IN PATIENTS WHO HAVE RECEIVED PRIOR ANTI-ANGIOGENIC THERAPY
- U-3240 USE OF VASCEPA TO REDUCE TRIGLYCERIDE LEVELS IN AN ADULT PATIENT HAVING TRIGLYCERIDE LEVELS OF AT LEAST ABOUT 500 MG/DL, ON ANTICOAGULANT/ANTIPLATELET/THROMBOLYTIC THERAPY, AND HAVING ATRIAL FIBRILLATION AND/OR ATRIAL FLUTTER
- U-3241 IN COMBINATION WITH ENDOCRINE THERAPY (TAMOXIFEN OR AN AROMATASE INHIBITOR) FOR THE ADJUVANT TREATMENT OF ADULT PATIENTS WITH HR-POSITIVE, HER2-NEGATIVE, NODE-POSITIVE, EARLY BREAST CANCER AT HIGH RISK OF RECURRENCE AND A KI-67 SCORE $\geq 20\%$
- U-3242 IN COMBINATION WITH FULVESTRANT FOR THE TREATMENT OF ADULT PATIENTS WITH HR-POSITIVE, HUMAN EPIDERMAL GROWTH FACTOR RECEPTOR 2 (HER2)-NEGATIVE ADVANCED OR METASTATIC BREAST CANCER WITH DISEASE PROGRESSION FOLLOWING ENDOCRINE THERAPY
- U-3243 IN COMBINATION WITH AN AROMATASE INHIBITOR AS INITIAL ENDOCRINE-BASED THERAPY FOR THE TREATMENT OF POSTMENOPAUSAL WOMEN, AND MEN WITH HR-POSITIVE, HUMAN EPIDERMAL GROWTH FACTOR RECEPTOR 2 (HER2)-NEGATIVE ADVANCED OR METASTATIC BREAST CANCER
- U-3244 A METHOD FOR TREATMENT OF PAIN IN ADULTS USING TRAMADOL HYDROCHLORIDE AND CELECOXIB
- U-3245 MAINTENANCE MONOTHERAPY TREATMENT OF BIPOLAR I DISORDER
- U-3246 FOR THE TREATMENT OF CHRONIC GRAFT VERSUS HOST DISEASE
- U-3247 FOR THE TREATMENT OF CHRONIC GRAFT VERSUS HOLD DISEASE
- U-3248 TREATING SECONDARY HYPERPARATHYROIDISM IN STAGE 3/4 CHRONIC KIDNEY DISEASE WITH SUSTAINED RELEASE 25-HYDROXYVITAMIN D TO REDUCE THE PATIENT'S SERUM PARATHYROID HORMONE LEVEL WHILE AVOIDING PTH OVERSUPPRESSION
- U-3249 TREATMENT OF CORONAVIRUS DISEASE 2019 (COVID-19) IN NON-HOSPITALIZED ADULTS AND PEDIATRIC PATIENTS (AT LEAST 12 YEARS OF AGE AND 40 KG)
- U-3250 METHOD OF TREATING PAIN, FOR EXAMPLE, TREATING POSTSURGICAL PAIN VIA INFILTRATION FOR LOCAL ANALGESIA OR VIA INTERSCALENE BRACHIAL PLEXUS NERVE BLOCK FOR REGIONAL ANALGESIA
- U-3251 ADMINISTRATION OF FERROUS BISGLYCINATE TABLETS
- U-3252 USE OF VUITY FOR THE TREATMENT OF PRESBYOPIA IN ADULTS
- U-3253 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 6 YEARS AND OLDER WHO HAVE IN THE CFTR GENE AT LEAST ONE F508DEL MUTATION OR A MUTATION THAT IS RESPONSIVE BASED ON IN VITRO DATA WITH A COMPOSITION ACCORDING TO AT LEAST ONE OF CLAIMS 1-9 OF US11179367
- U-3254 USE, IN COMBINATION WITH LOW-DOSE CYTARABINE, FOR THE TREATMENT OF NEWLY-DIAGNOSED ACUTE MYELOID LEUKEMIA (AML) IN ADULT PATIENTS WHO ARE ≥ 75 YEARS OLD OR WHO HAVE COMORBIDITIES THAT PRECLUDE USE OF INTENSIVE INDUCTION CHEMOTHERAPY
- U-3255 TREATMENT OF ADULTS WITH MODERATELY TO SEVERELY ACTIVE RHEUMATOID ARTHRITIS WHO HAVE HAD AN INADEQUATE RESPONSE OR INTOLERANCE TO ONE OR MORE TNF BLOCKERS
- U-3256 USE TO INCREASE LINEAR GROWTH IN PEDIATRIC PATIENTS WITH ACHONDROPLASIA WHO ARE 5 YEARS OF AGE AND OLDER WITH OPEN EPIPHYSES
- U-3257 TREATMENT OF TRD IN CONJUNCTION WITH AN ORAL ANTIDEPRESSANT BY NASALLY ADMINISTERING 56MG OR 84MG OF ESKETAMINE IN A MAINTENANCE PHASE WEEKLY OR 1X EVERY TWO WEEKS TO ADULTS WHO HAVE BEEN ADMINISTERED ESKETAMINE IN A INDUCTION PHASE FOR ABOUT 4 WEEKS
- U-3258 TREATMENT OF ADULT PATIENTS WITH LOCALLY ADVANCED UNRESECTABLE OR METASTATIC MALIGNANT PERIVASCULAR EPITHELIOID CELL TUMOR (PECOMA) WITH A DOSE BETWEEN ABOUT 56 MG/M2 AND ABOUT 100 MG/M2 ADMINISTERED ON DAYS 1 AND 8 OF A 21-DAY CYCLE
- U-3259 TREATMENT OF ADULT PATIENTS WITH LOCALLY ADVANCED UNRESECTABLE OR METASTATIC MALIGNANT PERIVASCULAR EPITHELIOID CELL TUMOR (PECOMA)
- U-3260 METHOD OF TREATING SPASTICITY
- U-3261 FOR CLEANSING OF THE COLON AS A PREPARATION FOR COLONOSCOPY
- U-3262 TREATING HYPOTENSION WITH ANGIOTENSIN II IN A PATIENT RECEIVING AN ANGIOTENSIN CONVERTING ENZYME INHIBITOR

PATENT AND EXCLUSIVITY TERMS

ADB 190 of 225

PATENT USE

- U-3263 METHOD FOR TREATING SPASTICITY
- U-3264 AS INITIAL ENDOCRINE-BASED THERAPY FOR THE TREATMENT OF ADULT PATIENTS WITH HORMONE RECEPTOR (HR)-POSITIVE, HUMAN EPIDERMAL GROWTH FACTOR RECEPTOR 2 (HER2)-NEGATIVE ADVANCED OR METASTATIC BREAST CANCER
- U-3265 IN COMBINATION WITH AN AROMATASE INHIBITOR AS INITIAL ENDOCRINE-BASED THERAPY FOR THE TREATMENT OF ADULT PATIENTS WITH HORMONE RECEPTOR (HR)-POSITIVE, HUMAN EPIDERMAL GROWTH FACTOR RECEPTOR 2 (HER2)-NEGATIVE ADVANCED OR METASTATIC BREAST CANCER
- U-3266 IN COMBINATION WITH FULVESTRANT AS INITIAL ENDOCRINE-BASED THERAPY OR FOLLOWING DISEASE PROGRESSION ON ENDOCRINE THERAPY IN POSTMENOPAUSAL WOMEN OR IN MEN, FOR THE TREATMENT OF HR-POSITIVE, HER2-NEGATIVE ADVANCED OR METASTATIC BREAST CANCER
- U-3267 USE OF LASTACFT TO TEMPORARY RELIEVE ITCHY EYES DUE TO POLLEN, RAGWEED, GRASS, ANIMAL HAIR AND DANDER
- U-3268 TREATMENT OF MULTIPLE SCLEROSIS IN PEDIATRIC PATIENTS 10 YEARS OF AGE AND OLDER AND WEIGHING LESS THAN OR EQUAL TO 40 KG
- U-3269 TREATMENT OF PRIMARY IMMUNOGLOBULIN A NEPHROPATHY (IGAN) IN ADULTS AT RISK OF RAPID DISEASE PROGRESSION
- U-3270 TREATMENT OF BACTERIAL VAGINOSIS IN FEMALE PATIENTS 12 YEARS OF AGE AND OLDER
- U-3271 TREATMENT OF BIPOLAR DEPRESSION
- U-3272 AS AN ADJUNCT TO DIET AND MAXIMALLY TOLERATED STATIN THERAPY FOR THE TREATMENT OF ADULTS WITH HETEROZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA (HEFH) OR CLINICAL ATHEROSCLEROTIC CARDIOVASCULAR DISEASE (ASCVD), BY INHIBITING EXPRESSION OF THE PCSK9 GENE
- U-3274 TREATMENT OF BIPOLAR I DISORDER, BIPOLAR II DISORDER, OR BIPOLAR DEPRESSION
- U-3275 TREATMENT OF ADULTS WITH ACTIVE PSORIATIC ARTHRITIS WHO HAVE HAD AN INADEQUATE RESPONSE OR INTOLERANCE TO ONE OR MORE TNF BLOCKERS
- U-3276 TREATMENT OF PATIENTS WITH PLAQUE PSORIASIS WHO ARE CANDIDATES FOR PHOTOTHERAPY OR SYSTEMIC THERAPY
- U-3277 USE FOR THE TREATMENT OF SEIZURES ASSOCIATED WITH TUBEROUS SCLEROSIS COMPLEX
- U-3278 A METHOD OF REDUCING POST-SURGICAL PAIN FOLLOWING OCULAR SURGERY
- U-3279 A METHOD OF TREATING POSTOPERATIVE INFLAMMATION FOLLOWING OCULAR SURGERY
- U-3280 METHOD OF TREATING ACNE VULGARIS WITH TOPICALLY APPLIED CORTEXOLONE 17ALPHA-PROPIONATE
- U-3281 TREATMENT OF SCHIZOPHRENIA IN ADULTS AND PEDIATRIC PATIENTS AGES 13 YEARS AND OLDER
- U-3282 DURING LEVOKETOCONAZOLE DOSAGE TITRATION FOR THE TREATMENT OF CUSHING'S SYNDROME IN PATIENTS WHO CONCOMITANTLY USE METFORMIN, MONITORING GLYCEMIA, KIDNEY FUNCTION AND VITAMIN B-12 AND ADJUSTING DOSAGE OF METFORMIN AS NEEDED
- U-3283 TREATMENT OF ENDOGENOUS HYPERCORTISOLEMIA IN PATIENTS WITH CUSHING'S SYNDROME FOR WHOM SURGERY IS NOT AN OPTION OR HAS NOT BEEN CURATIVE
- U-3284 PROPHYLAXIS OF THROMBOEMBOLIC DISEASES IN PEDIATRIC PATIENTS AGED 2 YEARS AND OLDER WITH CONGENITAL HEART DISEASE WHO HAVE UNDERGONE THE FONTAN PROCEDURE AND A BODY WEIGHT OF ≥ 50 KG
- U-3285 TREATMENT OF DVT AND/OR PE AND REDUCTION IN THE RISK OF RECURRENT DVT AND/OR PE IN PEDIATRIC PATIENTS (≥ 50 KG) ONCE DAILY WITH RAPID-RELEASE TABLET ADMINISTERED FOR AT LEAST 5 CONSECUTIVE DAYS AFTER AT LEAST 5 DAYS PARENTERAL ANTICOAGULANT TREATMENT
- U-3286 TREATMENT OF VENOUS THROMBOEMBOLISM (VTE) AND THE REDUCTION IN THE RISK OF RECURRENT VTE IN PEDIATRIC PATIENTS FROM BIRTH TO LESS THAN 18 YEARS WITH A BODY WEIGHT OF 30 KG TO 49.9 KG AFTER AT LEAST 5 DAYS OF INITIAL PARENTERAL ANTICOAGULANT TREATMENT
- U-3287 TREATMENT OF VENOUS THROMBOEMBOLISM (VTE) AND THE REDUCTION IN THE RISK OF RECURRENT VTE IN PEDIATRIC PATIENTS FROM BIRTH TO LESS THAN 18 YEARS WITH A BODY WEIGHT OF ≥ 50 KG AFTER AT LEAST 5 DAYS OF INITIAL PARENTERAL ANTICOAGULANT TREATMENT
- U-3288 PROPHYLAXIS OF PE, DVT AND/OR STROKE IN PEDIATRIC PATIENTS (≥ 50 KG) AGED 2 YEARS AND OLDER WITH CONGENITAL HEART DISEASE AFTER FONTAN PROCEDURE WITH ONCE DAILY, RAPID-RELEASE TABLET ADMINISTERED FOR AT LEAST 5 CONSECUTIVE DAYS

PATENT AND EXCLUSIVITY TERMS

ADB 191 of 225

PATENT USE

- U-3289 TREATMENT OF DVT AND/OR PE AND REDUCTION IN RISK OF RECURRENT DVT AND/OR PE IN PEDIATRIC PATIENTS (30-49.9 KG) ONCE DAILY WITH RAPID-RELEASE TABLET ADMINISTERED FOR AT LEAST 5 CONSECUTIVE DAYS AFTER AT LEAST 5 DAYS PARENTERAL ANTICOAGULANT TREATMENT
- U-3290 TREATMENT OF CHOLESTATIC PRURITUS IN PATIENTS WITH ALAGILLE SYNDROME (ALGS)
- U-3291 CYTALUX IS AN OPTICAL IMAGING AGENT INDICATED IN ADULT PATIENTS WITH OVARIAN CANCER AS AN ADJUNCT FOR INTRAOPERATIVE IDENTIFICATION OF MALIGNANT LESIONS
- U-3292 USE OF VASCEPA TO REDUCE TRIGLYCERIDE LEVELS IN AN ADULT PATIENT ON STATIN THERAPY AND HAVING ATRIAL FIBRILLATION OR ATRIAL FLUTTER AND TRIGLYCERIDE LEVELS OF ABOUT 500 MG/DL TO ABOUT 2,000 MG/DL
- U-3293 METHOD OF TREATING BACTERIAL VAGINOSIS BY SINGLE DOSE ADMINISTRATION OF A CLINDAMYCIN PHARMACEUTICAL GEL FORMULATION
- U-3294 METHOD OF TREATING A BACTERIAL INFECTION BY ADMINISTERING A RECONSTITUTED SOLID FORMULATION OF DAPTOMYCIN CONTAINING 31.0 TO 59.4% WT TOTAL MANNITOL AND SORBITOL
- U-3295 METHOD OF DELIVERING A COMBINATION OF OLOPATADINE HYDROCHLORIDE AND MOMETASONE FUROATE TO A NASAL AIRWAY
- U-3296 TREATMENT OF SEASONAL ALLERGIC RHINITIS BY NASALY ADMINISTERING A COMBINATION OF OLOPATADINE HYDROCHLORIDE AND MOMETASONE FUROATE TO A PEDIATRIC PATIENT
- U-3297 TREATMENT OF SEASONAL ALLERGIC RHINITIS BY NASALY ADMINISTERING A COMBINATION OF OLOPATADINE HYDROCHLORIDE AND MOMETASONE FUROATE
- U-3298 TREATMENT OF ADULTS AND PEDIATRIC PATIENTS 12 YEARS OF AGE AND OLDER WITH REFRACTORY, MODERATE TO SEVERE ATOPIC DERMATITIS WHOSE DISEASE IS NOT ADEQUATELY CONTROLLED WITH OTHER SYSTEMIC DRUG PRODUCTS, OR WHEN USE OF THOSE THERAPIES ARE INADVISABLE
- U-3299 TREATMENT OF ATTENTION DEFICIT HYPERACTIVITY DISORDER (ADHD) IN PEDIATRIC PATIENTS
- U-3300 PROPHYLAXIS TO PREVENT ATTACKS OF HEREDITARY ANGIOEDEMA (HAE) IN ADULTS AND PEDIATRIC PATIENTS 12 YEARS OF AGE AND OLDER
- U-3301 TREATMENT OF ADULT AND PEDIATRIC PATIENTS WITH HEPATIC VENO-OCCLUSIVE DISEASE (VOD), ALSO KNOWN AS SINUSOIDAL OBSTRUCTION SYNDROME (SOS), WITH RENAL OR PULMONARY DYSFUNCTION FOLLOWING HEMATOPOIETIC STEM-CELL TRANSPLANTATION (HSCT)
- U-3302 TREATMENT OF TRICHOMONIASIS IN PATIENTS 12 YEARS OF AGE AND OLDER
- U-3303 TOPICAL LESION-DIRECTED AND FIELD-DIRECTED TREATMENT OF ACTINIC KERATOSIS OF THE FACE AND SCALP WITH PHOTODYNAMIC THERAPY BY POSITIONING AN ILLUMINATION DEVICE IN AN APPROPRIATE DISTANCE AND ILLUMINATING THE TREATMENT AREA WITH NARROWBAND RED LIGHT
- U-3304 METHOD OF TREATING PARKINSON'S DISEASE BY ORALLY ADMINISTERING SEGMENTS OF A FUNCTIONALLY MULTISCORED, BILAYERED TABLET HAVING CARBIDOPA-25 MG/LEVODOPA-100 MG, EACH SEGMENT HAVING CARBIDOPA-6.25 MG/LEVODOPA-25 MG
- U-3305 METHOD OF TREATING PARKINSON'S DISEASE BY ORALLY ADMINISTERING A FUNCTIONALLY MULTISCORED, BILAYERED TABLET HAVING CARBIDOPA-25 MG/LEVODOPA-100 MG
- U-3306 TREATMENT OF ADULT PATIENTS WITH KRAS G12C-MUTATED LOCALLY ADVANCED OR METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC), AS DETERMINED BY AN FDA-APPROVED TEST, WHO HAVE RECEIVED AT LEAST ONE PRIOR SYSTEMIC THERAPY
- U-3307 TREATMENT OF HIV-1 INFECTION IN PEDIATRIC PATIENTS WEIGHING AT LEAST 35KG WITH NO PRIOR ANTIRETROVIRAL TREATMENT HISTORY OR TO REPLACE THE CURRENT ANTIRETROVIRAL REGIMEN IN THOSE WHO ARE VIROLOGICALLY SUPPRESSED
- U-3308 FOR USE IN COMBINATION WITH OTHER ANTIRETROVIRAL AGENTS TO TREAT HIV-1 INFECTION IN PEDIATRIC PATIENTS WEIGHING AT LEAST 35KG WITH NO PRIOR ANTIRETROVIRAL TREATMENT HISTORY OR TO REPLACE THE CURRENT ANTIRETROVIRAL REGIMEN IF VIROLOGICALLY SUPPRESSED
- U-3309 NORLIQVA IS INDICATED FOR THE TREATMENT OF HYPERTENSION, TO LOWER BLOOD PRESSURE IN ADULTS AND CHILDREN 6 YEARS OF AGE AND OLDER
- U-3310 NORLIQVA IS INDICATED FOR THE SYMPTOMATIC TREATMENT OF CHRONIC STABLE ANGINA
- U-3311 NORLIQVA IS INDICATED FOR THE TREATMENT OF CONFIRMED OR SUSPECTED VASOSPASTIC ANGINA
- U-3312 METHOD TO TREAT IDA IN ADULTS & PEDIATRIC PATIENTS 1 YR & OLDER WITH INTOLERANCE OR UNSATISFACTORY RESPONSE TO ORAL IRON & WEIGHING AT LEAST 40 KG BY

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

- INTRAVENOUSLY ADMINISTERING FERRIC CARBOXYMALTOSE TO PROVIDE AT LEAST ABOUT 0.6 G OF ELEMENTAL IRON
- U-3313 METHOD OF TREATING IDA IN ADULT PATIENTS WITH NON-DIALYSIS DEPENDENT CHRONIC KIDNEY DISEASE & WEIGHING AT LEAST 40 KG BY INTRAVENOUSLY ADMINISTERING FERRIC CARBOXYMALTOSE TO PROVIDE AT LEAST ABOUT 0.6 GRAMS OF ELEMENTAL IRON
- U-3314 METHOD OF TREATING IDA IN ADULTS & PEDIATRIC PATIENTS 1 YR & OLDER HAVING INTOLERANCE OR UNSATISFACTORY RESPONSE TO ORAL IRON ASSOCIATED WITH HEAVY UTERINE BLEEDING OR GASTROINTESTINAL DISORDER BY INTRAVENOUSLY ADMINISTERING FERRIC CARBOXYMALTOSE
- U-3315 METHOD TO TREAT IDA IN ADULTS & PEDIATRIC PATIENTS 1 YR & OLDER WITH INTOLERANCE OR UNSATISFACTORY RESPONSE TO ORAL IRON & WEIGHING AT LEAST 40 KG BY ADMINISTERING IV AT LEAST ABOUT 0.6 G OF IRON AS FERRIC CARBOXYMALTOSE IN ABOUT 15 MIN OR LESS
- U-3316 METHOD OF TREATING IDA IN ADULT PATIENTS WITH NON-DIALYSIS DEPENDENT CHRONIC KIDNEY DISEASE & WEIGHING AT LEAST 40 KG BY INTRAVENOUSLY ADMINISTERING AT LEAST ABOUT 0.6 GRAMS OF IRON AS FERRIC CARBOXYMALTOSE IN ABOUT 15 MINUTES OR LESS
- U-3317 POSITRON EMISSION TOMOGRAPHY (PET) OF PROSTATE-SPECIFIC MEMBRANE ANTIGEN (PSMA) POSITIVE LESIONS IN MEN WITH PROSTATE CANCER
- U-3318 MANAGEMENT OF MODERATE-TO-SEVERE PAIN BY INJECTION
- U-3319 METHOD OF USING A PYRUVATE KINASE ACTIVATOR FOR THE TREATMENT OF HEMOLYTIC ANEMIA IN ADULTS WITH PYRUVATE KINASE (PK) DEFICIENCY
- U-3320 METHOD FOR INCREASING THE LIFETIME OF RED BLOOD CELLS (RBCS) FOR THE TREATMENT OF HEMOLYTIC ANEMIA IN ADULTS WITH PYRUVATE KINASE (PK) DEFICIENCY
- U-3321 METHOD OF USING A PYRUVATE KINASE ACTIVATOR FOR THE TREATMENT OF HEMOLYTIC ANEMIA IN ADULTS WITH PYRUVATE KINASE (PK) DEFICIENCY BY ADMINISTERING A DAILY DOSE OF 10MG TO 100MG
- U-3322 USE FOR DETECTING NEUTRALIZING ANTIBODIES
- U-3323 METHOD OF REDUCING ADVERSE EFFECTS IN PATIENTS WHO ARE CONCOMITANTLY ADMINISTERED A SALT OF GAMMA-HYDROXYBUTYRATE AND DIVALPROEX SODIUM
- U-3324 METHOD OF TREATING PATIENTS WITH A SALT OF GAMMA-HYDROXYBUTYRATE WHEN DIVALPROEX SODIUM IS CONCOMITANTLY ADMINISTERED
- U-3325 METHOD FOR REDUCING THE RISK OF CARDIOVASCULAR DEATH AND HOSPITALIZATION FOR HEART FAILURE IN ADULTS WITH TYPE 2 DIABETES MELLITUS AND HEART FAILURE BY ONCE DAILY ADMINISTRATION OF EMPAGLIFLOZIN
- U-3326 A METHOD OF TREATING ANKYLOSING SPONDYLITIS BY ADMINISTERING THE FORMULATION OF TOFACITINIB OF CLAIMS 1, 8, 15-20, 27-31, 34-43, 45, 47, 49, 50, 54, 59, 63, 68-71, 73, 77, 82-84, AND 87-98
- U-3327 A METHOD OF TREATING PSORIATIC ARTHRITIS BY ADMINISTERING THE FORMULATION OF TOFACITINIB OF CLAIMS 1, 8, 15-20, 27-31, 34-44, 46, 48, 50, 53, 59, 62, 68-71, 73, 76, 82-84, AND 87-98
- U-3328 A METHOD OF TREATING RHEUMATOID ARTHRITIS BY ADMINISTERING THE FORMULATION OF TOFACITINIB OF CLAIMS 1, 2, 8, 9, 15-21, 27-31, 34-43, 50, 51, 59, 60, 68-71, 73, 74, 82-84 AND 87-98
- U-3329 A METHOD OF TREATING ULCERATIVE COLITIS, BY ADMINISTERING THE FORMULATION OF TOFACITINIB OF CLAIMS 1, 3, 8, 10, 15-20, 22, 27-31, 34-43, 50, 52, 59, 61, 68-71, 73, 75, 82-84 AND 87-98
- U-3330 TREATMENT OF GASTROINTESTINAL STROMAL TUMORS IN PATIENTS SUFFERING FROM PALMER-PLANTER ERYTHRODYSESTHESIA SYNDROME
- U-3331 TREATMENT OF MYELOFIBROSIS WITH PACRITINIB
- U-3332 USE OF PACRITINIB FOR INHIBITING JANUS ASSOCIATED KINASE 2 (JAK2)
- U-3333 ADJUVANT TREATMENT OF PATIENTS WITH GBRCA-MUTATED HUMAN EPIDERMAL GROWTH FACTOR RECEPTOR 2 (HER2)-NEGATIVE HIGH RISK EARLY BREAST CANCER WHO HAVE BEEN TREATED WITH NEOADJUVANT OR ADJUVANT CHEMOTHERAPY
- U-3334 A METHOD OF TRANSDERMAL DELIVERY OF DONEPEZIL FOR TREATING MILD, MODERATE AND SEVERE DEMENTIA OF THE ALZHEIMER'S TYPE
- U-3335 TREATMENT OF HOSPITAL-ACQUIRED BACTERIAL PNEUMONIA (HABP)
- U-3336 TREATMENT OF VENTILATOR-ASSOCIATED BACTERIAL PNEUMONIA (VABP)
- U-3337 ADMINISTERING DAILY A UNIT DOSAGE OF AN IRREVERSIBLE EGFR INHIBITOR COVALENTLY

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

- BINDING AS CLAIMED FOR 1ST LINE TREATMENT OF GEFITINIB OF ERLOTINIB RESISTANT METASTATIC NSCLC WITH EGFR EXON 19 DELETION OR EXON 21 L858R SUBSTITUTION
- U-3338 ADMINISTERING DAILY A UNIT DOSAGE OF AN IRREVERSIBLE EGFR INHIBITOR COVALENTLY BINDING AS CLAIMED FOR 1ST LINE TREATMENT OF GEFITINIB OR ERLOTINIB RESISTANT METASTATIC NSCLC WITH EGFR EXON 19 DELETION OR EXON 21 L858R SUBSTITUTION WITH T790M MUTATION
- U-3339 TREATMENT OF CF IN A PATIENT AGE 4 MONTHS TO <6 YEARS WHO HAS ONE MUTATION IN THE CFTR GENE THAT IS RESPONSIVE TO IVACAFTOR BASED ON CLINICAL AND/OR IN VITRO ASSAY DATA USING THE COMPOSITION RECITED IN CLAIM 1 OF US 11147770
- U-3340 A METHOD OF TREATING ATTENTION DEFICIT HYPERACTIVITY DISORDER BY ADMINISTERING A TRANSDERMAL COMPOSITION CONTAINING AMPHETAMINE
- U-3341 TREATMENT OF ADULTS WITH MODERATELY TO SEVERELY ACTIVE ULCERATIVE COLITIS WHO HAVE HAD AN INADEQUATE RESPONSE OR INTOLERANCE TO ONE OR MORE TNF BLOCKERS
- U-3342 TREATMENT OF NIGHTTIME SLEEP DISTURBANCES IN SMITH-MAGENIS SYNDROME BY AVOIDING THE ADMINISTRATION OF TASIMELTEON WITH BETA-ADRENERGIC RECEPTOR ANTAGONISTS
- U-3343 TREATMENT OF NON-24-HOUR SLEEP-WAKE DISORDER BY AVOIDING THE ADMINISTRATION OF TASIMELTEON WITH BETA-ADRENERGIC RECEPTOR ANTAGONISTS
- U-3344 A METHOD OF TREATING ADULTS WITH MULTIPLE MYELOMA USING DEXAMETHASONE IN COMBINATION WITH AN ANTI-MYELOMA PRODUCT
- U-3345 FOR TREATMENT OF ADULT PATIENTS WITH PROSTATE-SPECIFIC MEMBRANE ANTIGEN (PSMA)-POSITIVE METASTATIC CASTRATION-RESISTANT PROSTATE CANCER (MCRPC) WHO HAVE BEEN TREATED WITH ANDROGEN RECEPTOR (AR) PATHWAY INHIBITION AND TAXANE-BASED CHEMOTHERAPY
- U-3346 METHOD OF PROVIDING LOCAL OR REGIONAL ANALGESIA VIA INFILTRATION FOR LOCAL ANALGESIA OR VIA INTERSCALENE BRACHIAL PLEXUS NERVE BLOCK OR FEMORAL NERVE BLOCK FOR REGIONAL ANALGESIA
- U-3347 USE IN COMBINATION WITH CANNABIDIOL FOR THE TREATMENT OF SEIZURES ASSOCIATED WITH LENNOX-GASTAUT SYNDROME
- U-3348 TREATMENT OF HIV-1 INFECTION IN ADULTS AND ADOLESCENTS 12 YEARS OF AGE AND OLDER AND WEIGHING AT LEAST 35 KG
- U-3349 REINITIATION OF SCHIZOPHRENIA TREATMENT WHEREIN MORE THAN 6 MONTHS 3 WEEKS BUT LESS THAN 8 MONTHS HAVE ELAPSED SINCE THE LAST DOSE
- U-3350 ACUTE TREATMENT OF AGITATION ASSOCIATED WITH SCHIZOPHRENIA OR BIPOLAR I OR II DISORDER BY SUBLINGUAL ADMINISTRATION
- U-3351 METHOD OF TREATING IRON DEFICIENCY ANEMIA IN ADULT PATIENTS WITH NON-DIALYSIS DEPENDENT CHRONIC KIDNEY DISEASE BY INTRAVENOUSLY ADMINISTERING FERRIC CARBOXYMALTOSATE
- U-3352 METHOD TO TREAT IRON DEFICIENCY ANEMIA IN ADULTS & PEDIATRIC PATIENTS 1 YEAR AND OLDER WITH INTOLERANCE OR UNSATISFACTORY RESPONSE TO ORAL IRON BY INTRAVENOUSLY ADMINISTERING FERRIC CARBOXYMALTOSATE
- U-3353 TREATMENT IN COMBINATION WITH CABOTEGRAVIR OF HIV-1 INFECTION IN ADULTS AND ADOLESCENTS 12 AND OLDER TO REPLACE CURRENT REGIMEN IN THOSE WHO ARE VIROLOGICALLY SUPPRESSED ON A STABLE ANTIRETROVIRAL REGIMEN WITH NO HISTORY OF TREATMENT FAILURE
- U-3354 TREATMENT OF ADULT PATIENTS WITH PAROXYSMAL NOCTURNAL HEMOGLOBINURIA (PNH) BY ADMINISTRATION OF PEGCETACOPLAN SO AS TO REDUCE THE SENSITIVITY OF CELLS TO COMPLEMENT-DEPENDENT DAMAGE
- U-3355 AN ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS
- U-3356 TOPICAL TREATMENT OF INFLAMMATORY LESIONS OF ROSACEA IN ADULTS 18 YEARS OF AGE AND OLDER
- U-3357 TOPICAL TREATMENT OF INFLAMMATORY LESIONS OF ROSACEA IN ADULTS 65 YEARS OF AGE AND OLDER
- U-3358 USE OF VASCEPA TO REDUCE THE INCIDENCE OF MI IN AN ADULT PATIENT ON STATIN THERAPY AND WITH ELEVATED TRIGLYCERIDE LEVELS (>150 MG/DL), WHEREIN THE PATIENT EXPERIENCES ATRIAL FIBRILLATION AND/OR FLUTTER INSTEAD OF AN INCIDENCE OF MI
- U-3359 TREATMENT OF SCHIZOPHRENIA BY ADMINISTERING A DOSE UP TO TWO WEEKS BEFORE OR THREE WEEKS AFTER THE SCHEDULED SIX-MONTH DOSE
- U-3360 COMPLICATED INTRA-ABDOMINAL INFECTIONS (CIAI), USED IN COMBINATION WITH

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

- METRONIDAZOLE, IN ADULT AND PEDIATRIC PATIENTS (BIRTH TO LESS THAN 18 YEARS OF AGE)
- U-3361 COMPLICATED URINARY TRACT INFECTIONS (CUTI), INCLUDING PYELONEPHRITIS, IN ADULT AND PEDIATRIC PATIENTS (BIRTH TO LESS THAN 18 YEARS OLD)
- U-3362 TREATMENT OF BIPOLAR DEPRESSION MEDIATED BY THE 5-HT2A RECEPTOR, SEROTONIN TRANSPORTER, AND/OR DOPAMINE D1/D2 SIGNALING PATHWAYS
- U-3363 TREATMENT OF SCHIZOPHRENIA MEDIATED BY THE 5-HT2A RECEPTOR, SEROTONIN TRANSPORTER, AND/OR DOPAMINE D1/D2 SIGNALING PATHWAYS
- U-3364 TREATMENT OF BIPOLAR DISORDER I, BIPOLAR DISORDER II, OR BIPOLAR DEPRESSION
- U-3365 THE PRODUCT COMPOSITION (NATROBA) IS FOR THE TOPICAL TREATMENT OF HUMAN SCABIES MITE INFESTATIONS BY MELTING AND DELIVERING THE ACTIVE INGREDIENT, SPINOSAD, TO THE STRATUM CORNEUM WHERE SCABIES MITES LIVE AND BREED
- U-3366 VIVJOA IS INDICATED TO REDUCE THE INCIDENCE OF RECURRENT VULVOVAGINAL CANDIDIASIS (RVVC) IN FEMALES WITH A HISTORY OF RVVC WHO ARE NOT OF REPRODUCTIVE POTENTIAL
- U-3367 TREATMENT OF CORONAVIRUS DISEASE 2019 (COVID-19) IN ADULTS AND PEDIATRIC PATIENTS (AT LEAST 28 DAYS OF AGE AND 3 KG) REQUIRING HOSPITALIZATION
- U-3368 TREATMENT OF CORONAVIRUS DISEASE 2019 (COVID-19) IN NON-HOSPITALIZED ADULTS AND PEDIATRIC PATIENTS (AT LEAST 28 DAYS OF AGE AND 3 KG)
- U-3369 TREATMENT OF ADULTS AND PEDIATRIC PATIENTS 12 AND OLDER WITH SCLERODERMATOUS FORM OF CHRONIC GRAFT-VERSUS-HOST DISEASE (CGVHD) AFTER FAILURE OF AT LEAST TWO PRIOR LINES OF SYSTEMIC THERAPY
- U-3370 A METHOD FOR THE TREATMENT OF ADULT PATIENTS WITH STABLE WILSON'S DISEASE WHO ARE DE-COPPERED AND TOLERANT TO PENICILLAMINE
- U-3371 TREATMENT OF ADULTS WITH ACTIVE ANKYLOSING SPONDYLITIS WHO HAVE HAD AN INADEQUATE RESPONSE OR INTOLERANCE TO ONE OR MORE TNF BLOCKERS
- U-3372 TREATMENT OF CORONAVIRUS DISEASE 2019 (COVID-19) IN HOSPITALIZED ADULTS REQUIRING SUPPLEMENTAL OXYGEN, NON-INVASIVE OR INVASIVE MECHANICAL VENTILATION, OR EXTRACORPOREAL MEMBRANE OXYGENATION (ECMO)
- U-3373 TREATMENT OF ADULTS WITH SYMPTOMATIC NEW YORK HEART ASSOCIATION (NYHA) CLASS II-III OBSTRUCTIVE HYPERTROPHIC CARDIOMYOPATHY (HCM) TO IMPROVE FUNCTIONAL CAPACITY AND SYMPTOMS
- U-3374 TREATMENT OF SEIZURES ASSOCIATED WITH CYCLIN-DEPENDENT KINASE-LIKE 5 (CDKL5) DEFICIENCY DISORDER (CDD) IN PATIENTS 2 YEARS OF AGE AND OLDER
- U-3375 USE FOR THE TREATMENT OF ABSENCE SEIZURES IN PATIENTS WITH DRAVET SYNDROME
- U-3376 USE FOR THE TREATMENT OF ABSENCE SEIZURES IN PATIENTS WITH LENNOX-GASTAUT SYNDROME
- U-3377 TPOXX IS INDICATED FOR THE TREATMENT OF HUMAN SMALLPOX DISEASE IN ADULTS AND PEDIATRIC PATIENTS WEIGHING AT LEAST 3 KG
- U-3378 MOUNJARO IS INDICATED AS AN ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS
- U-3379 A METHOD FOR ADMINISTERING A THERAPEUTIC DOSE OF EPINEPHRINE
- U-3380 METHOD OF TREATING PAIN, FOR EXAMPLE, VIA INFILTRATION FOR LOCAL ANALGESIA OR VIA INTERSCALENE BRACHIAL PLEXUS NERVE BLOCK FOR REGIONAL ANALGESIA
- U-3381 TREATMENT OF GASTROINTESTINAL STROMAL TUMORS IN PATIENTS SUFFERING FROM HYPERTENSION
- U-3382 TREATMENT OF GASTROINTESTINAL STROMAL TUMORS IN PATIENTS SUFFERING FROM A GRADE 3 ADVERSE REACTION WHILE BEING ADMINISTERED RIPRETINIB DAILY
- U-3383 TREATING NEWLY DIAGNOSED AML CHARACTERIZED BY THE PRESENCE OF A MUTANT ALLELE OF IDH1 BY ADMINISTERING A ONCE DAILY 500 MG ORAL DOSE OF IVOSIDENIB TO A SUBJECT THAT HAS NOT INGESTED A HIGH-FAT MEAL, IN COMBINATION WITH AZACITIDINE
- U-3384 A METHOD FOR TREATING AML BY ADMINISTERING A PHARMACEUTICAL COMPOSITION COMPRISING IVOSIDENIB WHEREIN THE AML IS NEWLY DIAGNOSED AND CHARACTERIZED BY A MUTANT IDH1 AND THE COMPOSITION IS ADMINISTERED IN COMBINATION WITH AZACITIDINE
- U-3385 A METHOD FOR TREATING NEWLY DIAGNOSED AML WITH IVOSIDENIB AND AZACITIDINE WHEREIN THE AML HAS AN IDH1 MUTATION CAPABLE OF CONVERTING ALPHA-KETOGLUTARATE TO 2-HYDROXYGLUTARATE (2HG)
- U-3386 A METHOD FOR TREATING A CANCER CHARACTERIZED BY AN IDH1 MUTATION WITH IVOSIDENIB

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

- IN COMBINATION WITH AZACITIDINE WHEREIN THE CANCER IS NEWLY DIAGNOSED AML
- U-3387 A METHOD FOR TREATING NEWLY DIAGNOSED AML CHARACTERIZED BY THE PRESENCE OF A MUTANT ALLELE OF IDH1 SELECTED FROM R132H, R132C, R132L, R132V, R132S AND R132GF BY ADMINISTERING IVOSIDENIB AND AZACITIDINE
- U-3388 USE OF ELAGOLIX 200 MG BID FOR 6 MONTHS TO MANAGE MODERATE TO SEVERE PAIN ASSOCIATED WITH ENDOMETRIOSIS IN PREMENOPAUSAL WOMEN TO REDUCE DYSMENORRHEA AND NON-MENSTRUAL PELVIC PAIN
- U-3389 USE OF ELAGOLIX 200 MG BID FOR 6 MONTHS TO MANAGE MODERATE TO SEVERE PAIN ASSOCIATED WITH ENDOMETRIOSIS IN PREMENOPAUSAL WOMEN HAVING DYSpareunia ASSOCIATED WITH ENDOMETRIOSIS
- U-3390 SODIUM THIOSULFATE INJECTION IS ADMINISTERED BY INTRAVENOUS INJECTION
- U-3391 A METHOD OF TITRATING AN OPIOID TO MANAGE PAIN SEVERE ENOUGH TO REQUIRE DAILY, AROUND-THE-CLOCK, LONG-TERM OPIOID TREATMENT AND FOR WHICH ALTERNATIVE TREATMENTS ARE INADEQUATE
- U-3392 A METHOD OF TITRATING AN OPIOID TO MANAGE NEUROPATHIC PAIN ASSOCIATED WITH DIABETIC PERIPHERAL NEUROPATHY SEVERE ENOUGH TO REQUIRE DAILY, AROUND-THE-CLOCK, LONG-TERM OPIOID TREATMENT AND FOR WHICH ALTERNATIVE TREATMENT OPTIONS ARE INADEQUATE
- U-3393 METHOD OF TREATING ACUTE PAIN DUE TO MINOR STRAINS, SPRAINS, AND CONTUSIONS USING A DICLOFENAC PATCH CONTAINING HEPARIN FOR ONCE DAILY ADMINISTRATION WHERE HEPARIN IS NOT RELEASED
- U-3394 SODIUM NITRITE INJECTION IS ADMINISTERED BY INTRAVENOUS INJECTION
- U-3395 SODIUM NITRITE INJECTION IS INDICATED FOR SEQUENTIAL USE WITH SODIUM THIOSULFATE FOR THE TREATMENT OF ACUTE CYANIDE POISONING
- U-3396 AMVUTTRA IS INDICATED FOR THE TREATMENT OF THE POLYNEUROPATHY OF HEREDITARY TRANSTHYRETIN-MEDIATED AMYLOIDOSIS IN ADULTS
- U-3397 TADLIQ IS INDICATED FOR THE TREATMENT OF PULMONARY ARTERIAL HYPERTENSION (PAH) (WHO GROUP 1) TO IMPROVE EXERCISE ABILITY
- U-3398 FOR CHRONIC WEIGHT MANAGEMENT IN ADULTS WITH BMI ≥ 30 KG/M² OR BMI ≥ 27 KG/M² WITH A WEIGHT-RELATED COMORBIDITY, AND PATIENTS AGE 12-17 WITH BMI ≥ 25 KG/M² IN THE 95TH PERCENTILE OR GREATER (STANDARDIZED FOR AGE AND SEX)
- U-3399 FOR CHRONIC WEIGHT MANAGEMENT IN ADULTS WITH BMI ≥ 30 KG/M², AND PATIENTS AGE 12-17 WITH BMI ≥ 30 KG/M² AND IN THE 95TH PERCENTILE OR GREATER (STANDARDIZED FOR AGE AND SEX), EACH HAVING A WEIGHT-RELATED COMORBIDITY
- U-3400 FOR USE AFTER RADIOLABELING WITH GALLIUM-68, FOR POSITRON EMISSION TOMOGRAPHY OF PROSTATE-SPECIFIC MEMBRANE ANTIGEN (PSMA)-POSITIVE LESIONS IN MEN WITH PROSTATE CANCER
- U-3401 CAROSPIR IS INDICATED AS AN ADD-ON THERAPY FOR THE TREATMENT OF HYPERTENSION, TO LOWER BLOOD PRESSURE IN ADULT PATIENTS WHO ARE NOT ADEQUATELY CONTROLLED ON OTHER AGENTS
- U-3402 CAROSPIR IS INDICATED FOR THE MANAGEMENT OF EDEMA IN ADULT CIRRHOTIC PATIENTS WHEN EDEMA IS NOT RESPONSIVE TO FLUID AND SODIUM RESTRICTION
- U-3403 METHOD OF REVERSING OR INHIBITING THE PROGRESS OF UNRESECTABLE, RECURRENT, OR REFRACTORY INFLAMMATORY MYOFIBROBLASTIC TUMOR (IMT) THAT IS ALK-POSITIVE IN ADULT AND PEDIATRIC PATIENTS 1 YEAR OF AGE AND OLDER
- U-3404 FOR THE TOPICAL TREATMENT OF NONSEGMENTAL VITILIGO IN ADULT AND PEDIATRIC PATIENTS 12 YEARS OF AGE AND OLDER
- U-3405 TREATMENT OF HIV-1 INFECTION IN ADULTS AND ADOLESCENTS 12 YEARS OF AGE AND OLDER AND WEIGHING AT LEAST 35 KG BY MONTHLY ADMINISTRATION OF RILPIVIRINE SUSPENSION AS PART OF COMBINATION THERAPY
- U-3406 USE OF FENFLURAMINE AT REDUCED AMOUNTS WITH STIRIPENTOL FOR THE TREATMENT OF SEIZURES ASSOCIATED WITH DRAVET SYNDROME
- U-3407 USE OF FENFLURAMINE AT REDUCED AMOUNTS WITH STIRIPENTOL FOR THE TREATMENT OF SEIZURES ASSOCIATED WITH LENNOX GASTAUT SYNDROME
- U-3408 TOPICAL TREATMENT OF PLAQUE PSORIASIS, INCLUDING INTERTRIGINOUS AREAS, IN PATIENTS 12 YEARS OF AGE AND OLDER. (1)
- U-3409 TREATING ADULT PATIENTS WITH INTERMEDIATE-2 OR HIGH-RISK PRIMARY OR SECONDARY MYELOFIBROSIS, MONITORING THIAMINE LEVELS AND ADMINISTERING THIAMINE OR A THIAMINE EQUIVALENT

PATENT AND EXCLUSIVITY TERMS

ADB 196 of 225

PATENT USE

- U-3410 A METHOD OF INJECTING AN IMPLANT
- U-3411 TREATING NON-EARLY SPMS BY ADMINISTERING ORAL CLADRIBINE AT A FIXED DOSE PER PATIENT, PER BODY WEIGHT AND PER TREATMENT YEAR, WHICH FIXED DOSE IS 1.75 +/- 0.2 MG/KG, TO BE ADMINISTERED WITHIN MONTHS 1 AND 2 IN EACH OF 2 ADJACENT TREATMENT YEARS
- U-3412 TREATMENT OF ADULTS WITH RELAPSED, REFRACTORY OR PREVIOUSLY UNTREATED CHRONIC LYMPHOCYTIC LEUKEMIA INCLUDING A DOSE RAMP-UP AND IN COMBINATION WITH OBINUTUZUMAB IN MULTIPLE 28-DAY DOSING CYCLES FOLLOWED BY ADMINISTRATION IN ABSENCE OF OBINUTUZUMAB
- U-3413 INDICATED AS ADJUNCTIVE THERAPY FOR THE TREATMENT OF PARTIAL-ONSET SEIZURES, PRIMARY GENERALIZED TONIC-CLONIC SEIZURES, AND SEIZURES ASSOCIATED WITH LENNOX-GASTAUT SYNDROME IN PATIENTS 2 YEARS OF AGE AND OLDER
- U-3414 INDICATED AS INITIAL MONOTHERAPY FOR THE TREATMENT OF PARTIAL-ONSET OR PRIMARY GENERALIZED TONIC-CLONIC SEIZURES IN PATIENTS 2 YEARS OF AGE AND OLDER
- U-3415 INDICATED FOR THE PREVENTIVE TREATMENT OF MIGRAINE IN PATIENTS 12 YEARS AND OLDER
- U-3416 TREATMENT OF GASTROINTESTINAL STROMAL TUMORS IN PATIENTS BEING TREATED CONCURRENTLY WITH A CYP3A4 INHIBITOR
- U-3417 TREATMENT OF POSTSURGICAL PAIN PROVIDING ANALGESIA TO A PATIENT FOR UP TO 72 HOURS, FOR EXAMPLE, AFTER FOOT AND ANKLE, SMALL-TO-MEDIUM OPEN ABDOMINAL, AND LOWER EXTREMITY TOTAL JOINT ARTHROPLASTY SURGICAL PROCEDURES VIA INSTILLATION
- U-3418 A METHOD OF LOADING MEDICATION INTO A SYRINGE AND DELIVERING THE MEDICATION TO A TREATMENT SITE
- U-3419 DEXTROMETHORPHAN AND BUPROPION IN COMBINATION TO TREAT MAJOR DEPRESSIVE DISORDER
- U-3420 A METHOD OF TREATING TESTOSTERONE DEFICIENCY IN MEN
- U-3421 TREATMENT OF COMPLICATED URINARY TRACT INFECTIONS (CUTI) INCLUDING PYELONEPHRITIS CAUSED BY THE FOLLOWING SUSCEPTIBLE MICROORGANISMS: ESCHERICHIA COLI, KLEBSIELLA PNEUMONIA, ENTEROBACTER CLOACAE SPECIES COMPLEX WITH MEROPENEM & VABORBACTAM AS SPECIFIED
- U-3422 TREATMENT OF PEDIATRIC PATIENTS AGE 1 YEAR AND OLDER WITH CHRONIC GRAFT-VERSUS-HOST DISEASE (CGVHD) AFTER FAILURE OF ONE OR MORE LINES OF SYSTEMIC THERAPY
- U-3423 METHOD OF TREATING GASTROINTESTINAL STROMAL TUMORS IN PATIENTS SUFFERING FROM GRADE 2 OR GRADE 3 MYALGIA WHILE BEING ADMINISTERED RIPRETINIB DAILY
- U-3424 TREATMENT OF CYSTIC FIBROSIS IN A PATIENT AGE 1-5 YEARS WHO IS HOMOZYGOUS FOR THE F508DEL MUTATION IN THE CFTR GENE USING LUMACAFTOR AND IVACAFTOR
- U-3425 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 6 YEARS AND OLDER WHO HAVE IN THE CFTR GENE AT LEAST ONE F508DEL MUTATION OR A MUTATION THAT IS RESPONSIVE BASED ON IN VITRO DATA WITH A COMPOUND OF CLAIM 1 OR COMPOSITION OF CLAIM 29 OF US11426407
- U-3426 TREATMENT OF CYSTIC FIBROSIS IN A PATIENT AGE 1-5 YEARS WHO IS HOMOZYGOUS FOR THE F508DEL MUTATION IN THE CFTR GENE USING THE DOSAGE UNIT COMPRISING LUMACAFTOR AS RECITED IN CLAIM 1 OF US PATENT 8716338 AND IVACAFTOR
- U-3427 TREATMENT OF CYSTIC FIBROSIS IN A PATIENT AGE 1-5 YEARS WHO IS HOMOZYGOUS FOR THE F508DEL MUTATION IN THE CFTR GENE USING LUMACAFTOR FORM I AND IVACAFTOR
- U-3428 TREATMENT OF CYSTIC FIBROSIS IN A PATIENT AGE 1-5 YEARS WHO IS HOMOZYGOUS FOR THE F508DEL MUTATION IN THE CFTR GENE USING THE DOSAGE UNIT COMPRISING LUMACAFTOR AND IVACAFTOR AS RECITED IN CLAIM 1 OF US PATENT 9192606
- U-3429 TREATMENT OF CYSTIC FIBROSIS IN A PATIENT AGE 1-5 YEARS WHO IS HOMOZYGOUS FOR THE F508DEL MUTATION IN THE CFTR GENE USING LUMACAFTOR AND A SOLID COMPOSITION COMPRISING AMORPHOUS AND LESS THAN ABOUT 30% CRYSTALLINE IVACAFTOR
- U-3430 TREATMENT OF CYSTIC FIBROSIS IN A PATIENT AGE 1-5 YEARS WHO IS HOMOZYGOUS FOR THE F508DEL MUTATION IN THE CFTR GENE USING A PHARMACEUTICAL COMPOSITION ACCORDING TO CLAIM 2 OF U.S. PATENT NO. 10,597,384, FURTHER COMPRISING IVACAFTOR
- U-3431 TREATMENT OF NOCTURIA DUE TO NOCTURNAL POLYURIA IN ADULTS WHO AWAKEN AT LEAST 2 TIMES PER NIGHT TO VOID BY INDUCING AN ANTIDIURETIC EFFECT BY INTRANASALLY ADMINISTERING A PLUME OF DROPLETS COMPRISING A DOSE OF ABOUT 0.05-5 MCG DESMOPRESSIN
- U-3432 TREATMENT OF CATAPLEXY OR EXCESSIVE DAYTIME SLEEPINESS (EDS) IN PATIENTS 7 YEARS OF AGE AND OLDER WITH NARCOLEPSY WITH A MIXTURE OF SODIUM, POTASSIUM, MAGNESIUM, AND CALCIUM SALTS OF GHB ADMINISTERED BETWEEN 2 AND 4 HOURS AFTER EATING

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

- U-3433 INDICATED FOR THE TREATMENT OF SPASTICITY RESULTING FROM MULTIPLE SCLEROSIS
- U-3434 TREATMENT OF MODERATE-TO-SEVERE PLAQUE PSORIASIS IN ADULTS WHO ARE CANDIDATES FOR SYSTEMIC THERAPY OR PHOTOTHERAPY
- U-3435 METHOD OF TREATING IDA IN ADULT PATIENTS WITH NON-DIALYSIS DEPENDENT CHRONIC KIDNEY DISEASE & WEIGHING AT LEAST 46.7 KG BY INTRAVENOUSLY ADMINISTERING FERRIC CARBOXYMALTOSE TO PROVIDE AT LEAST 0.7 G OF ELEMENTAL IRON IN 15 MINUTES
- U-3436 METHOD OF TREATING IDA IN ADULT PATIENTS WITH NON-DIALYSIS DEPENDENT CHRONIC KIDNEY DISEASE & WEIGHING AT LEAST 46.7 KG BY INTRAVENOUSLY ADMINISTERING FERRIC CARBOXYMALTOSE TO PROVIDE AT LEAST 0.7 G OF ELEMENTAL IRON IN LESS THAN 15 MINUTES
- U-3437 METHOD TO TREAT IDA IN ADULTS & PEDIATRIC PATIENTS 1 YEAR & OLDER WITH INTOLERANCE OR UNSATISFACTORY RESPONSE TO ORAL IRON & HEAVY UTERINE BLEEDING OR GI DISORDER BY ADMINISTERING IV FERRIC CARBOXYMALTOSE TO GIVE AT LEAST 0.7 G OF IRON IN 15 MINUTES
- U-3438 METHOD TO TREAT IDA IN ADULTS & PEDIATRIC PATIENTS 1 YR & OLDER WITH INTOLERANCE OR UNSATISFACTORY RESPONSE TO ORAL IRON & WEIGHING AT LEAST 46.7 KG BY ADMINISTERING IV FERRIC CARBOXYMALTOSE TO PROVIDE AT LEAST 0.7 G OF ELEMENTAL IRON IN < 15 MINUTES
- U-3439 METHOD OF TREATING PAIN, FOR EXAMPLE, VIA INFILTRATION FOR LOCAL ANALGESIA OR VIA NERVE BLOCK, FOR EXAMPLE, NTERSCALENE BRACHIAL PLEXUS FOR REGIONAL ANALGESIA
- U-3440 A METHOD OF ADMINISTERING APREPITANT FOR PREVENTION OF POST-OPERATIVE NAUSEA AND VOMITING
- U-3441 A METHOD OF TREATING CYSTINURIA BY ORALLY ADMINISTERING TIOPRONIN WITH FOOD TO PREVENT CYSTINE STONE FORMATION IN ADULTS AND PEDIATRIC PATIENTS WITH SEVERE HOMOZYGOUS CYSTINURIA
- U-3442 TREATMENT OF ADVANCED GASTROINTESTINAL STROMAL TUMOR IN PATIENTS PREVIOUSLY ADMINISTERED AT LEAST THREE TYROSINE KINASE INHIBITORS, WHERE ONE OF THE KINASE INHIBITORS IS IMATINIB
- U-3443 A METHOD OF REDUCING OTOTOXICITY IN A HUMAN PEDIATRIC PATIENT ABOUT 5 YEARS OF AGE OR UNDER WITH LOCALIZED MEDULLOBLASTOMA COMPRISING ADMINISTERING SODIUM THIOSULFATE ABOUT SIX HOURS AFTER ADMINISTRATION OF CISPLATIN
- U-3444 TREATMENT OF DEPRESSIVE SYMPTOMS IN ADULTS WITH MDD WITH ACUTE SUICIDAL IDEATION OR BEHAVIOR BY NASALLY ADMINISTERING 56MG OR 84MG OF ESKETAMINE AS A PHARMACEUTICAL COMPOSITION TWICE PER WEEK IN CONJUNCTION WITH AN ORAL ANTIDEPRESSANT
- U-3445 TREATMENT OF TREATMENT-RESISTANT DEPRESSION IN ADULTS BY NASALLY ADMINISTERING 56MG OR 84MG OF ESKETAMINE TWICE PER WEEK AS A PHARMACEUTICAL COMPOSITION IN CONJUNCTION WITH AN ORAL ANTIDEPRESSANT
- U-3446 TREATMENT OF TREATMENT-RESISTANT DEPRESSION IN ADULTS BY NASALLY ADMINISTERING 56MG OR 84MG OF ESKETAMINE AS A PHARMACEUTICAL COMPOSITION IN CONJUNCTION WITH AN ORAL ANTIDEPRESSANT
- U-3447 A METHOD OF TREATING CORONARY ARTERY DISEASE
- U-3448 A METHOD OF TREATING HYPERTENSION
- U-3449 USE IN COMBINATION WITH LENALIDOMIDE PLUS DEXAMETHASONE FOR THE TREATMENT OF PATIENTS WITH RELAPSED OR REFRACTORY MULTIPLE MYELOMA WHO HAVE RECEIVED ONE TO THREE LINES OF THERAPY
- U-3450 TREATMENT OF ADULT PATIENTS WITH LOCALLY ADVANCED OR METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) WITH A REARRANGED DURING TRANSFECTION (RET) GENE FUSION, AS DETECTED BY AN FDA-APPROVED TEST
- U-3451 TREATMENT OF ADULT AND PEDIATRIC PATIENTS 12 YEARS OF AGE AND OLDER WITH ADVANCED OR METASTATIC MEDULLARY THYROID CANCER (MTC) WITH A RET MUTATION, AS DETECTED BY AN FDA-APPROVED TEST, WHO REQUIRE SYSTEMIC THERAPY
- U-3452 TREATMENT OF ADULT AND PEDIATRIC PATIENTS 12 YEARS OF AGE AND OLDER WITH ADVANCED OR METASTATIC THYROID CANCER WITH A RET GENE FUSION, AS DETECTED BY AN FDA-APPROVED TEST, WHO REQUIRE SYSTEMIC THERAPY AND WHO ARE RADIOACTIVE IODINE-REFRACTORY
- U-3453 TREATMENT OF ADULT PATIENTS WITH LOCALLY ADVANCED OR METASTATIC SOLID TUMORS WITH A RET GENE FUSION THAT HAVE PROGRESSED ON OR FOLLOWING PRIOR SYSTEMIC TREATMENT OR WHO HAVE NO SATISFACTORY ALTERNATIVE TREATMENT OPTIONS
- U-3454 METHOD OF TREATING OPEN-ANGLE GLAUCOMA OR OCULAR HYPERTENSION IN PATIENTS

PATENT AND EXCLUSIVITY TERMS

ADB 198 of 225

PATENT USE

U-3455 TREATMENT OF OCULAR INFLAMMATION AND PAIN FOLLOWING OPHTHALMIC SURGERY

U-3456 METHOD OF TREATING INTRAHEPATIC CHOLANGIOCARCINOMA

U-3457 METHOD OF INDUCING OCULAR ANESTHESIA

U-3458 A METHOD OF TREATING SEIZURES

U-3459 TREATING SICKLE CELL DISEASE BY ADMINISTERING VOXELOTOR AND ANOTHER ACTIVE AGENT

U-3460 TREATMENT OF AMYOTROPHIC LATERAL SCLEROSIS (ALS) IN ADULTS

U-3461 INDICATED TO RAISE BLOOD PRESSURE IN ADULT PATIENTS WITH SEVERE, ACUTE HYPOTENSION

U-3462 USE OF A LIQUID FORMULATION COMPRISING FUROSEMIDE TO TREAT CONGESTION DUE TO FLUID OVERLOAD (EDEMA) IN ADULTS WITH NYHA CLASS II/III CHRONIC HEART FAILURE

U-3463 USE OF TRINTELLIX FOR THE TREATMENT OF MAJOR DEPRESSIVE DISORDER (MDD) IN ADULTS

U-3464 TREATMENT OF PREVIOUSLY TREATED, UNRESECTABLE LOCALLY ADVANCED OR METASTATIC CHOLANGIOCARCINOMA WITH A FGFR2 FUSION OR OTHER REARRANGEMENT BY ADMINISTERING ONCE DAILY A TABLET CONTAINING ABOUT 0.5 MG TO ABOUT 10 MG OF PEMIGATINIB

U-3465 TREATMENT OF PREVIOUSLY TREATED, UNRESECTABLE LOCALLY ADVANCED OR METASTATIC CHOLANGIOCARCINOMA WITH A FGFR2 FUSION OR OTHER REARRANGEMENT BY ADMINISTERING PEMIGATINIB QD FOR 14 DAYS THEN NOT ADMINISTERING PEMIGATINIB FOR 7 DAYS IN A 21-DAY CYCLE

U-3466 TREATMENT OF PREVIOUSLY TREATED, UNRESECTABLE LOCALLY ADVANCED OR METASTATIC CHOLANGIOCARCINOMA WITH A FGFR2 FUSION OR OTHER REARRANGEMENT BY ADMINISTERING PEMIGATINIB IN A DAILY DOSE OF ABOUT 5 MG TO ABOUT 20 MG

U-3467 PREVENTION AND TREATMENT OF POST-OPERATIVE NAUSEA AND VOMITING

U-3468 TREATMENT OF AMYOTROPHIC LATERAL SCLEROSIS

U-3469 AN ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS

U-3470 METHOD OF TREATING COMPLICATED URINARY TRACT INFECTIONS (CUTI), INCLUDING PYELONEPHRITIS, COMPRISING ADMINISTERING CEFIDEROCOL SULFATE TOSYLATE

U-3471 METHOD OF TREATING HOSPITAL-ACQUIRED BACTERIAL PNEUMONIA AND VENTILATOR-ASSOCIATED BACTERIAL PNEUMONIA (HABP/VABP) COMPRISING ADMINISTERING CEFIDEROCOL SULFATE TOSYLATE

U-3472 METHOD TO TREAT IDA IN ADULTS WITH INTOLERANCE OR UNSATISFACTORY RESPONSE TO ORAL IRON & HEAVY UTERINE BLEEDING OR GI DISORDER WEIGHING AT LEAST 40 KG BY ADMINISTERING IV FERRIC CARBOXYMALTOSE TO GIVE AT LEAST 0.6 G OF IRON IN 15 MINUTES OR LESS

U-3473 METHOD TO TREAT IRON DEFICIENCY ANEMIA IN ADULTS WITH INTOLERANCE OR UNSATISFACTORY RESPONSE TO ORAL IRON & WEIGHING AT LEAST 40 KG BY INTRAVENOUSLY ADMINISTERING FERRIC CARBOXYMALTOSE TO PROVIDE AT LEAST 0.6 G OF ELEMENTAL IRON IN 15 MINUTES OR LESS

U-3474 METHOD TO TREAT IRON DEFICIENCY ANEMIA IN ADULTS WITH NON-DIALYSIS DEPENDENT CHRONIC KIDNEY DISEASE & WEIGHING AT LEAST 40 KG BY INTRAVENOUSLY ADMINISTERING FERRIC CARBOXYMALTOSE TO PROVIDE AT LEAST 0.6 G OF ELEMENTAL IRON IN 15 MINUTES OR LESS

U-3475 REDUCTION OF THE FREQUENCY OF ABDOMINAL PAIN AND DIARRHEA, IN AN INFLAMMATORY BOWEL DISEASE WITH DIARRHEA (IBS-D) PATIENT, WITH ELUXADOLINE TWICE DAILY WITH FOOD

U-3476 TREATMENT OF ADULTS WITH MODERATE HEPATIC IMPAIRMENT AND RELAPSED OR REFRACTORY ADVANCED RENAL CELL CARCINOMA FOLLOWING TWO OR MORE PRIOR SYSTEMIC ANTI-CANCER THERAPIES WITH 1MG TIVOZANIB HCL ORALLY FOR 21 DAYS FOLLOWED BY NO DRUG FOR 7 DAYS

U-3477 ACCELERATE THE TRANSIT OF A BARIUM MEAL THROUGH THE SMALL BOWEL, THEREBY DECREASING THE TIME AND EXTENT OF RADIATION ASSOCIATED WITH FLUOROSCOPY AND X-RAY EXAMINATION OF THE INTESTINAL TRACT

U-3478 STIMULATE GALLBLADDER CONTRACTION, AS MAY BE ASSESSED BY VARIOUS METHODS OF DIAGNOSTIC IMAGING, OR TO OBTAIN BY DUODENAL ASPIRATION A SAMPLE OF CONCENTRATED BILE FOR ANALYSIS

U-3479 STIMULATE PANCREATIC SECRETION IN COMBINATION WITH SECRETIN PRIOR TO OBTAINING A DUODENAL ASPIRATE FOR ANALYSIS

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

- U-3480 STIMULATE GALLBLADDER CONTRACTION. AS MAY BE ASSESSED BY VARIOUS METHODS OF DIAGNOSTIC IMAGING, OR TO OBTAIN BY DUODENAL ASPIRATION A SAMPLE OF CONCENTRATED BILE FOR ANALYSIS
- U-3481 DIAGNOSIS OF GALL BLADDER DISORDERS OR OTHER DIAGNOSTIC IMAGING BY STIMULATING GALLBLADDER CONTRACTION, AS MAY BE ASSESSED BY VARIOUS METHODS OF DIAGNOSTIC IMAGING, OR TO OBTAIN BY DUODENAL ASPIRATION A SAMPLE OF CONCENTRATED BILE FOR ANALYSIS
- U-3482 DIAGNOSIS OF PANCREATIC DISORDERS BY STIMULATING PANCREATIC SECRETION IN COMBINATION WITH SECRETIN PRIOR TO OBTAINING A DUODENAL ASPIRATE FOR ANALYSIS
- U-3483 DIAGNOSTIC IMAGING BY ACCELERATING THE TRAN IT OFA BARIUM MEAL THROUGH THE SMALL BOWEL, THEREBY DECREASING THE TIME AND EXTENT OF RADIATION ASSOCIATED WITH FLUOROSCOPY AND X-RAY EXAMINATION OF THE INTESTINAL TRACT
- U-3484 TREATMENT OF CORONAVIRUS DISEASE 2019 (COVID-19) IN ADULTS AND PEDIATRIC PATIENTS (AT LEAST 28 DAYS OF AGE AND 3 KG) REQUIRING HOSPITALIZATION AND FOR WHOM CONCOMITANT USE OF CHLOROQUINE, OR AN ANALOG OR SALT THEREOF, IS NOT RECOMMENDED
- U-3485 TREATMENT OF CORONAVIRUS DISEASE 2019 (COVID-19) IN NON-HOSPITALIZED ADULTS AND PEDIATRIC PATIENTS (AT LEAST 28 DAYS OF AGE AND 3 KG) AND FOR WHOM CONCOMITANT USE OF CHLOROQUINE, OR AN ANALOG OR SALT THEREOF, IS NOT RECOMMENDED
- U-3486 TREATMENT OF ADULT PATIENTS WITH SMALL LYMPHOCYTIC LYMPHOMA
- U-3487 TREATMENT OF ADULTS WITH ACTIVE NON-RADIOGRAPHIC AXIAL SPONDYLOARTHRITIS WITH OBJECTIVE SIGNS OF INFLAMMATION WHO HAVE HAD AN INADEQUATE RESPONSE OR INTOLERANCE TO ONE OR MORE TNF BLOCKERS
- U-3488 TREATMENT OF SPASTICITY RESULTING FROM MULTIPLE SCLEROSIS, PARTICULARLY FOR THE RELIEF OF FLEXOR SPASMS AND CONCOMITANT PAIN, CLONUS, AND MUSCULAR RIGIDITY
- U-3489 TREATMENT OF SPASTICITY RESULTING FROM SPINAL CORD INJURIES AND OTHER SPINAL CORD DISEASES
- U-3490 TREATMENT OF ADULT PATIENTS WITH KRAS G12C-MUTATED LOCALLY ADVANCED OR METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC), AS DETERMINED BY AN FDA APPROVED TEST, WHO HAVE RECEIVED AT LEAST ONE PRIOR SYSTEMIC THERAPY
- U-3491 METHOD OF TREATING LYMPHOMA
- U-3492 METHOD OF TREATING SARCOMA
- U-3493 THE TREATMENT OF RELAPSING FORMS OF MULTIPLE SCLEROSIS (MS) IN PATIENTS 10 YEARS OF AGE AND OLDER
- U-3494 ONCE DAILY TOPICAL TREATMENT OF PERSISTENT FACIAL ERYTHEMA ASSOCIATED WITH ROSACEA IN FEMALE ADULTS
- U-3495 A METHOD OF TREATING ACUTE MYELOID LEUKEMIA (AML) IN PATIENTS WITH AN ISOCITRATE DEHYDROGENASE-1 (IDH1) MUTATION
- U-3496 A METHOD OF TREATING A CANCER CHARACTERIZED BY AN IDH1 MUTATION WHERE THE CANCER IS ACUTE MYELOID LEUKEMIA (AML)
- U-3497 A METHOD OF TREATING A CANCER WHERE THE CANCER IS ACUTE MYELOID LEUKEMIA (AML)
- U-3498 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 6 YEARS AND OLDER WHO HAVE IN THE CFTR GENE AT LEAST ONE F508DEL MUTATION OR A MUTATION THAT IS RESPONSIVE BASED ON IN VITRO DATA BY ADMINISTERING DAILY ELX (200 MG OR 100 MG); TEZ; AND IVA
- U-3499 TREATMENT OF ACUTE BACTERIAL SKIN AND SKIN STRUCTURE INFECTIONS (ABSSSI) IN ADULT AND PEDIATRIC PATIENTS USING A TWO-DOSE REGIMEN OF DALBAVANCIN
- U-3500 TREATMENT OF ADULT PATIENTS WITH SEVERE ALOPECIA AREATA
- U-3501 PALBOCICLIB FOR HR-POS. HER2-NEG. ADVANCED OR METASTATIC BREAST CANCER IN COMBO WITH AN AROMATASE INHIBITOR IN PTS AS INITIAL ENDOCRINE-BASED THERAPY OR WITH FULVESTRANT WITH DISEASE PROGRESSION AFTER ENDOCRINE THERAPY
- U-3502 TREATMENT OF A UREA CYCLE DISORDER INVOLVING DEFICIENCIES OF CARBAMYLPHOSPHATE SYNTHETASE, ORNITHINE TRANSCARBAMYLASE, OR ARGININOSUCCINIC ACID SYNTHETASE
- U-3503 ADJUNCTIVE THERAPY TO ANTIDEPRESSANTS FOR THE TREATMENT OF MAJOR DEPRESSIVE DISORDER
- U-3504 A METHOD TO DECREASE THE INCIDENCE OF CHEMOTHERAPY-INDUCED MYELOSUPPRESSION IN ADULT PATIENTS WHEN ADMINISTERED PRIOR TO A PLATINUM/ETOPOSIDE-CONTAINING REGIMEN THAT INCLUDES AN IMMUNE CHECKPOINT INHIBITOR FOR EXTENSIVE-STAGE SMALL CELL CANCER

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

- U-3505 FOR TOPICAL TREATMENT OF MODERATE AD IN NON-IMMUNOCOMPROMISED PATIENTS, WITH BASELINE BSA OF 3-20% AND ITCH NRS SCORE OF ≥ 4 , WHOSE DISEASE IS NOT ADEQUATELY CONTROLLED WITH TOPICAL PRESCRIPTION THERAPIES OR WHEN THOSE THERAPIES ARE NOT ADVISABLE
- U-3506 TREATMENT OF INDOLENT SYSTEMIC MASTOCYTOSIS (ISM)
- U-3507 IN COMBINATION WITH OTHER ANTIRETROVIRAL(S), FOR THE TREATMENT OF HIV-1 INFECTION IN HEAVILY-TREATMENT EXPERIENCED ADULTS WITH MULTIDRUG RESISTANT HIV-1 INFECTION
- U-3508 REDUCTION IN THE INCIDENCE OF RECURRENT VULVOVAGINAL CANDIDIASIS (RVVC) IN ADULT AND POST-MENARCHAL PEDIATRIC FEMALES
- U-3509 AS-NEEDED TREATMENT OR PREVENTION OF BRONCHOCONSTRICTION AND REDUCTION OF THE RISK OF EXACERBATIONS IN PATIENTS WITH ASTHMA 18 YEARS OF AGE AND OLDER
- U-3510 COMBINATION TREATMENT OF COLORECTAL CANCER INCLUDING RAS WILD-TYPE HER2 (ERBB2)-POSITIVE OR -OVEREXPRESSING UNRESECTABLE OR METASTATIC COLORECTAL CANCER
- U-3511 TREATMENT OF IRRITABLE BOWEL SYNDROME WITH DIARRHEA (IBS-D) IN ADULT FEMALE SUBJECTS AND SYMPTOMS THEREOF
- U-3512 TREATMENT OF IRRITABLE BOWEL SYNDROME WITH DIARRHEA (IBS-D) IN ADULT FEMALE SUBJECTS
- U-3513 TREATMENT OF SCHIZOPHRENIA IN ADULTS
- U-3514 INCREASING BLOOD PRESSURE IN A PATIENT HAVING DISTRIBUTIVE SHOCK
- U-3515 A METHOD FOR ADMINISTERING A THERAPEUTIC DOSE OF AN OPIOID ANTAGONIST COMPRISING AN EMERGENCY SYRINGE DEVICE
- U-3516 A METHOD FOR ADMINISTERING A THERAPEUTIC DOSE OF AN OPIOID ANTAGONIST COMPRISING AN EMERGENCY SYRINGE DEVICE INCLUDING A NEEDLE GUARD
- U-3517 A METHOD FOR ADMINISTERING A THERAPEUTIC DOSE OF AN OPIOID ANTAGONIST COMPRISING AN EMERGENCY SYRINGE DEVICE INCLUDING A WINDOW CONFIGURED TO ALLOW THE USER TO VIEW THE OPIOID ANTAGONIST IN THE SYRINGE
- U-3518 TREATMENT OF ADULT PATIENTS WITH RELAPSED OR REFRACTORY MANTLE CELL LYMPHOMA (MCL) AFTER AT LEAST TWO LINES OF SYSTEMIC THERAPY, INCLUDING A BTK INHIBITOR
- U-3519 TREATMENT WITH LENVIMA BY ADMINISTERING LENVIMA AS A SUSPENSION
- U-3520 TREATMENT OF EXCESSIVE DAYTIME SLEEPINESS BY ADMINISTERING A COMPOSITION COMPRISING SOLRIAMFETOL HYDROCHLORIDE AND 2-CHLOROPROPANE, WHEREIN THE COMPOSITION COMPRISES LESS THAN ABOUT 5 PPM 2CHLOROPROPANE
- U-3521 TREATMENT OF EXCESSIVE DAYTIME SLEEPINESS BY ADMINISTERING SOLRIAMFETOL TO A SUBJECT HAVING MILD, MODERATE, OR SEVERE RENAL IMPAIRMENT
- U-3522 METHOD OF TREATING EMESIS
- U-3523 TREATMENT OF AN ER-POSITIVE BREAST CANCER
- U-3524 TREATMENT OF AN ER-POSITIVE BREAST CANCER FOLLOWING AT LEAST ONE LINE OF ENDOCRINE THERAPY
- U-3525 TREATMENT OF CF IN PATIENTS AGED 6 YEARS AND OLDER WHO HAVE IN THE CFTR GENE AT LEAST ONE F508DEL MUTATION OR A MUTATION THAT IS RESPONSIVE BASED ON IN VITRO DATA BY ADMINISTERING THE COMPOSITION RECITED IN US 11564916 CLAIM 1
- U-3526 TREATMENT OF CF IN A PATIENT AGE 1 TO <6 YEARS AND WEIGHING 7 KG OR MORE WHO IS HOMOZYGOUS FOR THE F508DEL MUTATION IN THE CFTR GENE USING THE COMPOSITION RECITED IN CLAIM 1 OF US 11564916
- U-3527 TREATMENT OF CF IN A PATIENT AGE 6 YEARS AND OLDER WHO IS HOMOZYGOUS FOR F508DEL OR HAS AT LEAST ONE CFTR GENE MUTATION RESPONSIVE TO TEZ/IVA BASED ON IN VITRO DATA AND/OR CLINICAL EVIDENCE USING THE COMPOSITION RECITED IN US 11564916 CLAIM 1
- U-3528 TREATMENT OF CF IN A PATIENT AGE 4 MONTHS TO <6 YEARS WHO HAS ONE MUTATION IN THE CFTR GENE THAT IS RESPONSIVE TO IVACAFTOR BASED ON CLINICAL AND/OR IN VITRO ASSAY DATA USING THE COMPOSITION RECITED IN CLAIM 1 OF US 11564916
- U-3529 TREATMENT OF CF IN A PATIENT AGE 6 YEARS AND OLDER WHO IS HOMOZYGOUS FOR THE F508DEL MUTATION IN THE CFTR GENE USING THE COMPOSITION RECITED IN CLAIM 1 OF US 11564916
- U-3530 TREATMENT OF CF IN A PATIENT AGE 6 YEARS AND OLDER WHO HAS ONE MUTATION IN THE CFTR GENE THAT IS RESPONSIVE TO IVACAFTOR BASED ON CLINICAL AND/OR IN VITRO ASSAY DATA USING THE COMPOSITION RECITED IN CLAIM 1 OF US 11564916

PATENT AND EXCLUSIVITY TERMS

ADB 201 of 225

PATENT USE

- U-3531 TREATMENT OF A TYPE 2 DIABETES PATIENT WITH INSUFFICIENT GLYCEMIC CONTROL DESPITE METFORMIN THERAPY USING A COMPOSITION COMPRISING AN EXTENDED RELEASE CORE COMPRISING METFORMIN AND AN OUTER COATING COMPRISING EMPAGLIFLOZIN AND LINAGLIPTIN
- U-3532 METHOD FOR REDUCING THE RISK OF CARDIOVASCULAR DEATH AND HOSPITALIZATION FOR HEART FAILURE IN ADULTS WITH TYPE 2 DIABETES MELLITUS BY ONCE DAILY ADMINISTRATION OF EMPAGLIFLOZIN
- U-3533 METHOD FOR REDUCING THE RISK OF CARDIOVASCULAR DEATH IN ADULT PATIENTS WITH TYPE 2 DIABETES MELLITUS AND CARDIOVASCULAR DISEASE BY ONCE DAILY ADMINISTRATION OF 10 MG OR 25 MG OF EMPAGLIFLOZIN
- U-3534 PREVENTIVE TREATMENT OF MIGRAINE IN ADULTS
- U-3535 A METHOD OF TREATING ANEMIA
- U-3536 TREATMENT OF DEPRESSION IN ADULTS WITH MOD AND ACUTE SUICIDAL IDEATION OR BEHAVIOR IN CONJUNCTION WITH AN ORAL ANTIDEPRESSANT BY NASALLY ADMINISTERING 56MG OR 84MG OF ESKETAMINE IN A MAINTENANCE PHASE WEEKLY OR LX EVERY 2 WEEKS AFTER INDUCTION PHASE
- U-3537 TREATMENT OF GASTROINTESTINAL STROMAL TUMORS IN PATIENTS SUFFERING FROM GRADE 2 OR GRADE 3 ARTHRALGIA WHILE BEING ADMINISTERED RIPRETINIB DAILY
- U-3538 METHOD OF TREATING ATTENTION DEFICIT HYPERACTIVITY DISORDER
- U-3539 TREATING ACQUIRED, GENERALIZED HYPOACTIVE SEXUAL DESIRE DISORDER (HSDD) IN A PREMENOPAUSAL FEMALE PATIENT WITH CONTROLLED HYPERTENSION BY INJECTING BREMELANOTIDE MORE THAN ONCE WITH AT LEAST 24 HOURS BETWEEN DOSES AND NO MORE THAN 8 DOSES PER MONTH
- U-3540 TREATMENT OF GEOGRAPHIC ATROPHY SECONDARY TO AGE-RELATED MACULAR DEGENERATION BY INTRAVITREAL ADMINISTRATION OF PEGCETACOPLAN
- U-3541 TREATMENT OF GEOGRAPHIC ATROPHY SECONDARY TO AGE-RELATED MACULAR DEGENERATION BY ADMINISTERING COMPLEMENT INHIBITOR PEGCETACOPLAN
- U-3542 TREATMENT OF GEOGRAPHIC ATROPHY SECONDARY TO AGE-RELATED MACULAR DEGENERATION BY INTRAVITREAL ADMINISTRATION OF PEGCETACOPLAN AND ALSO ADMINISTERING AN ANTI-VEGF AGENT
- U-3543 TREATMENT TO INCREASE BONE DENSITY IN MEN WITH OSTEOPOROSIS AT HIGH RISK FOR FRACTURE
- U-3544 TREATMENT OF CF IN PATIENTS AGED 6 YEARS AND OLDER WHO HAVE IN THE CFTR GENE AT LEAST ONE F508DEL MUTATION OR A RESPONSIVE MUTATION BASED ON IN VITRO DATA BY ADMINISTERING ELEXACAFTOR, IVACAFTOR, AND A SOLID DISPERSION OF TEZACAFTOR AND A POLYMER
- U-3545 TREATMENT OF CF IN A PATIENT AGE 6 YEARS AND OLDER WHO IS HOMOZYGOUS FOR F508DEL OR HAS AT LEAST ONE CFTR GENE MUTATION RESPONSIVE TO TEZ/IVA BASED ON IN VITRO DATA AND/OR CLINICAL EVIDENCE USING THE COMPOSITION RECITED IN US 11578062 CLAIM 6 OR 13
- U-3546 IN COMBINATION WITH ENDOCRINE THERAPY (TAMOXIFEN OR AN AROMATASE INHIBITOR) FOR THE ADJUVANT TREATMENT OF ADULT PATIENTS WITH HR-POSITIVE, HER2-NEGATIVE, NODE POSITIVE, EARLY BREAST CANCER AT HIGH RISK OF RECURRENCE
- U-3547 INTRAVENOUS SOTALOL DOSING REGIMEN FOR ACHIEVING STEADY STATE EXPOSURE IN A FACILITY THAT CAN PROVIDE ELECTROCARDIOGRAPHIC MONITORING
- U-3548 TREATMENT OF AML BY ORALLY ADMINISTERING VENETOCLAX WITH AZACITIDINE OR DECITABINE OR LOW-DOSE CYTARABINE TO ADULTS 75 YEARS OR OLDER OR HAVING CERTAIN COMORBIDITIES PER A DOSE RAMP-UP INCLUDING AN INITIAL 100 MG OR A FINAL 400 MG PER DAY DOSE
- U-3549 INTRAVENOUS SOTALOL DOSING REGIMEN FOR USE IN A FACILITY THAT CAN PROVIDE ELECTROCARDIOGRAPHIC MONITORING
- U-3550 FOR THE TOPICAL TREATMENT OF NONSEGMENTAL VITILIGO IN ADULT AND PEDIATRIC PATIENTS 12 YEARS OF AGE AND OLDER, IN THE ABSENCE OF LASER OR PHOTOTHERAPY, WHEREIN THE VITILIGO AFFECTS AT LEAST ONE OF THE LOWER EXTREMITIES, TRUNK, AND FEET OF THE PATIENT
- U-3551 FOR THE TOPICAL TREATMENT OF NONSEGMENTAL VITILIGO IN ADULT AND PEDIATRIC PATIENTS 12 YEARS OF AGE AND OLDER, IN THE ABSENCE OF PHOTOTHERAPY, WHEREIN THE VITILIGO AFFECTS AT LEAST ONE OF THE LOWER EXTREMITIES, TRUNK, AND FEET OF THE PATIENT
- U-3552 TREATMENT OF FRIEDREICH'S ATAXIA IN ADULTS AND ADOLESCENTS AGED 16 YEARS AND

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

OLDER

- U-3553 A METHOD OF TREATING MILD TO MODERATE ACUTE PAIN IN ADULTS BY ADMINISTERING 975-1000 MG OF ACETAMINOPHEN AND 292.5-300 MG OF IBUPROFEN IN A SINGLE ADMINISTRATION
- U-3554 AS A SINGLE AGENT FOR THE TREATMENT OF ADULT PATIENTS WITH HISTIOCYTIC NEOPLASMS
- U-3555 ADMINISTRATION OF ZAVEGEPANT FOR ACUTE TREATMENT OF MIGRAINE WITH OR WITHOUT AURA
- U-3556 TREATMENT OF RETT SYNDROME OR A SYMPTOM THEREOF
- U-3557 TREATMENT OF PARKINSON'S DISEASE, POST-ENCEPHALITIC PARKINSONISM, AND SYMPTOMATIC PARKINSONISM THAT MAY FOLLOW CARBON MONOXIDE INTOXICATION OR MANGANESE INTOXICATION
- U-3558 AN ADJUNCTIVE TREATMENT OF ADULT PATIENTS WITH TAVNEOS (AVACOPAN) WITH SEVERE ACTIVE ANCA-ASSOCIATED VASCULITIS (GPA AND MPA) IN COMBINATION WITH STANDARD THERAPY INCLUDING GLUCOCORTICOIDS
- U-3559 A METHOD OF TARGETING RELEASE OF A NON-STEROIDAL ANTI-INFLAMMATORY DRUG (NSAID) TO THE SMALL INTESTINE OF THE SUBJECT WHEN ADMINISTERED ORALLY
- U-3560 FOR THE TREATMENT OF VERNAL KERATOCONJUNCTIVITIS IN CHILDREN AND ADULTS
- U-3561 TREATMENT OF PRESBYOPIA IN ADULTS BY ADMINISTRATION OF PILOCARPINE HCl FORMULATION ONCE DAILY
- U-3562 TREATMENT OF PRESBYOPIA IN ADULTS BY ADMINISTRATION OF PILOCARPINE HCl FORMULATION TWICE DAILY
- U-3563 DEXTROMETHORPHAN AND BUPROPION IN COMBINATION TO INCREASE DEXTROMETHORPHAN PLASMA LEVELS
- U-3564 MEKINIST IS INDICATED, IN COMBINATION WITH DABRAFENIB, FOR THE TREATMENT OF PEDIATRIC PATIENTS 1 YEAR OF AGE AND OLDER WITH LOW-GRADE GLIOMA (LGG) WITH A BRAF V600E MUTATION WHO REQUIRE SYSTEMIC THERAPY
- U-3565 TAFINLAR IS INDICATED, IN COMBINATION WITH TRAMETINIB, FOR THE TREATMENT OF PEDIATRIC PATIENTS 1 YEAR OF AGE AND OLDER WITH LOW-GRADE GLIOMA (LGG) WITH A BRAF V600E MUTATION WHO REQUIRE SYSTEMIC THERAPY
- U-3566 TREATMENT OF CANDIDEMIA AND INVASIVE CANDIDIASIS WITH REZAFUNGIN BY INTRAVENOUS ADMINISTRATION
- U-3567 TO INCREASE MEAN ARTERIAL BLOOD PRESSURE IN ADULT PATIENTS WITH HYPOTENSION ASSOCIATED WITH SEPTIC SHOCK
- U-3568 TREATMENT OF PREVIOUSLY TREATED, UNRESECTABLE LOCALLY ADVANCED OR METASTATIC CHOLANGIOCARCINOMA WITH A FGFR2 FUSION OR OTHER REARRANGEMENT BY ADMINISTERING PEMIGATINIB WHILE AVOIDING THE CONCOMITANT USE OF STRONG AND MODERATE CYP3A INHIBITORS
- U-3569 TREATMENT OF PREVIOUSLY TREATED, UNRESECTABLE LOCALLY ADVANCED OR METASTATIC CHOLANGIOCARCINOMA WITH A FGFR2 FUSION OR OTHER REARRANGEMENT BY ADMINISTERING PEMIGATINIB WHILE AVOIDING THE CONCOMITANT USE OF STRONG AND MODERATE CYP3A INDUCERS
- U-3570 TREATMENT OF RELAPSED OR REFRACTORY MYELOID/LYMPHOID NEOPLASMS WITH FGFR1 REARRANGEMENT BY ADMINISTERING PEMIGATINIB WHILE AVOIDING THE CONCOMITANT USE OF STRONG AND MODERATE CYP3A INDUCERS
- U-3571 TREATMENT OF RELAPSED OR REFRACTORY MYELOID/LYMPHOID NEOPLASMS WITH FGFR1 REARRANGEMENT BY ADMINISTERING PEMIGATINIB WHILE AVOIDING THE CONCOMITANT USE OF STRONG AND MODERATE CYP3A INHIBITORS
- U-3572 RAISE FOLATE LEVELS IN WOMEN WHO CHOOSE TO USE AN ORAL CONTRACEPTIVE AS THEIR METHOD OF CONTRACEPTION FOR THE PURPOSE OF REDUCING THE RISK OF A NEURAL TUBE DEFECT IN A PREGNANCY
- U-3573 TREATMENT OF MODERATE ACNE VULGARIS IN WOMEN AT LEAST 14 YEARS OF AGE IF THE PATIENT DESIRES AN ORAL CONTRACEPTIVE FOR BIRTH CONTROL
- U-3574 TREATMENT OF MOOD CHANGES AND/OR ANXIETY AS SYMPTOMS OF PREMENSTRUAL DYSPHORIC DISORDER (PMDD) IN WOMEN WHO CHOOSE TO USE AN ORAL CONTRACEPTIVE AS THEIR METHOD OF CONTRACEPTION
- U-3575 TREATMENT OF AMYOTROPHIC LATERAL SCLEROSIS (ALS) IN ADULTS WHO HAVE A MUTATION IN THE SUPEROXIDE DISMUTASE 1 (SOD1) GENE
- U-3576 TREATMENT OF NARCOLEPSY-RELATED CATAPLEXY OR EXCESSIVE DAYTIME SLEEPINESS USING A ONCE-DAILY PHARMACEUTICAL FORMULATION COMPRISING AN OXYBATE

PATENT AND EXCLUSIVITY TERMS

ADB 203 of 225

PATENT USE

- U-3577 TREATMENT OF NARCOLEPSY-RELATED CATAPLEXY OR EXCESSIVE DAYTIME SLEEPINESS USING A SINGLE DAILY, BEDTIME DOSE OF A GAMMA-HYDROXYBUTYRATE FORMULATION
- U-3578 TREATMENT OF NARCOLEPSY, CATAPLEXY OR EXCESSIVE DAYTIME SLEEPINESS USING A ONCE-NIGHTLY GAMMA-HYDROXYBUTYRATE FORMULATION
- U-3579 TREATMENT OF NARCOLEPSY AND ASSOCIATED DISORDERS AND SYMPTOMS USING A COMPOSITION COMPRISING GAMMA-HYDROXYBUTYRATE ONCE DAILY
- U-3580 TREATMENT OF NARCOLEPSY USING A DOSE PROPORTIONAL, ORAL PHARMACEUTICAL COMPOSITION COMPRISING GAMMA-HYDROXYBUTYRATE ONCE DAILY
- U-3581 TREATMENT OF CATAPLEXY OR EXCESSIVE DAYTIME SLEEPINESS IN ADULTS WITH NARCOLEPSY
- U-3582 LIQREV IS INDICATED FOR THE TREATMENT OF PULMONARY ARTERIAL HYPERTENSION (WHO GROUP 1) IN ADULTS TO IMPROVE EXERCISE ABILITY AND DELAY CLINICAL WORSENING
- U-3583 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 2 TO <6 YEARS WHO HAVE IN THE CFTR GENE AT LEAST ONE F508DEL MUTATION OR A RESPONSIVE MUTATION BASED ON IN VITRO DATA WITH AN EFFECTIVE AMOUNT OF ELX, TEZ, AND IVA
- U-3584 TREATMENT OF CF IN PATIENTS AGED 2 TO <6 YEARS WHO HAVE IN THE CFTR GENE AT LEAST ONE F508DEL MUTATION OR A RESPONSIVE MUTATION BASED ON IN VITRO DATA BY ADMINISTERING ELEXACAFOTOR, IVACAFOTOR, AND A SOLID DISPERSION OF TEZACAFOTOR AND A POLYMER
- U-3585 TREATMENT OF CF IN PATIENTS AGED 2 TO <6 YEARS WHO HAVE IN THE CFTR GENE AT LEAST ONE F508DEL MUTATION OR A MUTATION THAT IS RESPONSIVE BASED ON IN VITRO DATA COMPRISING ADMINISTERING THE COMPOSITION RECITED IN US 11564916 CLAIM 1
- U-3586 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 2 TO <6 YEARS AND OLDER WHO HAVE IN THE CFTR GENE AT LEAST ONE F508DEL MUTATION OR A MUTATION THAT IS RESPONSIVE BASED ON IN VITRO DATA BY ADMINISTERING DAILY ELX (100 MG OR 80 MG); TEZ; AND IVA
- U-3587 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 6 YEARS AND OLDER WHO HAVE IN THE CFTR GENE AT LEAST ONE F508DEL MUTATION OR A RESPONSIVE MUTATION BASED ON IN VITRO DATA WITH AN EFFECTIVE AMOUNT OF ELX, TEZ, AND IVA
- U-3588 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 2 TO <6 YEARS AND OLDER WHO HAVE AT LEAST ONE F508DEL MUTATION IN THE CFTR GENE OR A MUTATION IN THE CFTR GENE THAT IS RESPONSIVE BASED ON IN VITRO DATA WITH ELEXACAFOTOR, TEZACAFOTOR, AND IVACAFOTOR
- U-3589 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 2 TO <6 YEARS WHO HAVE AT LEAST ONE F508DEL MUTATION IN THE CFTR GENE OR A MUTATION IN THE CFTR GENE THAT IS RESPONSIVE BASED ON IN VITRO DATA WITH ELEXACAFOTOR, TEZACAFOTOR, AND IVACAFOTOR
- U-3590 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 2 TO <6 YEARS WHO HAVE AT LEAST ONE F508DEL MUTATION IN THE CFTR GENE OR A MUTATION IN THE CFTR GENE THAT IS RESPONSIVE BASED ON IN VITRO DATA WITH AN EFFECTIVE AMOUNT OF ELX, TEZ, AND IVA
- U-3591 TREATMENT OF CF IN PATIENTS AGED 2 TO <6 YEARS WHO HAVE AT LEAST ONE F508DEL MUTATION OR A MUTATION IN THE CFTR GENE THAT IS RESPONSIVE BASED ON IN VITRO DATA USING A SOLID COMPOSITION COMPRISING ELX, TEZ, AMORPHOUS IVA, AND < ~30% CRYSTALLINE IVA
- U-3592 TREATMENT OF A MODERATE TO MILD CLINICAL PHENOTYPE OF CF IN PATIENTS AGED 2 TO <6 YEARS WHO HAVE AT LEAST ONE F508DEL MUTATION IN THE CFTR GENE OR A MUTATION IN THE CFTR GENE THAT IS RESPONSIVE BASED ON IN VITRO DATA WITH ELX, TEZ, AND IVA
- U-3593 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 2 TO <6 YEARS WHO HAVE A R117H MUTATION IN THE CFTR GENE WITH ELEXACAFOTOR, TEZACAFOTOR, AND IVACAFOTOR
- U-3594 TREATMENT OF IRON DEFICIENCY ANEMIA (DIA) IN ADULTS PATIENTS WHO HAVE INTOLERANCE TO ORAL IRON OR HAVE HAD UNSATISFACTORY RESPONSE TO ORAL IRON, WHO HAVE NON-HEMODIALYSIS DEPENDENT CHRONIC KIDNEY DISEASE, BY ADMINISTERING FERRIC DERISOMALTOSE
- U-3595 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 2 TO <6 YEARS WHO HAVE IN THE CFTR GENE AT LEAST ONE F508DEL MUTATION OR A MUTATION THAT IS RESPONSIVE BASED ON IN VITRO DATA WITH A COMPOUND OF CLAIM 1 OR COMPOSITION OF CLAIM 29 OF US 11426407
- U-3596 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 2 TO <6 YEARS WHO HAVE AT LEAST ONE F508DEL MUTATION IN THE CFTR GENE WITH AN EFFECTIVE AMOUNT OF A PHARMACEUTICAL COMPOSITION COMPRISING ELEXACAFOTOR, TEZACAFOTOR, AND IVACAFOTOR
- U-3597 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 2 TO <6 YEARS WHO HAVE IN THE CFTR GENE AT LEAST ONE F508DEL MUTATION OR A MUTATION THAT IS RESPONSIVE BASED ON IN VITRO DATA WITH A COMPOSITION ACCORDING TO ANY ONE OF CLAIMS 1-3 AND 7-9 OF

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

US11179367

- U-3598 TREATMENT OF CF IN PATIENTS AGED 2 TO <6 YEARS WHO HAVE AT LEAST ONE F508DEL MUTATION OR A MUTATION IN THE CFTR GENE THAT IS RESPONSIVE BASED ON IN VITRO DATA COMPRISING ADMINISTERING A COMPOSITION ACCORDING TO CLAIM 1 OF US 11147770
- U-3599 TREATMENT OF CF IN PATIENTS AGED 2 TO <6 YEARS WHO HAVE AT LEAST ONE F508DEL MUTATION OR A MUTATION IN THE CFTR GENE THAT IS RESPONSIVE BASED ON IN VITRO DATA COMPRISING ADMINISTERING A COMPOSITION ACCORDING TO CLAIM 1 OF US 10272046
- U-3600 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 2 TO <6 YEARS WHO HAVE AT LEAST ONE F508DEL MUTATION IN THE CFTR GENE OR A MUTATION IN THE CFTR GENE THAT IS RESPONSIVE BASED ON IN VITRO DATA WITH A COMPOSITION ACCORDING TO CLAIM 1 OF US 10081621
- U-3601 TREATMENT OF A DISORDER TREATABLE WITH GAMMA-HYDROXYBUTYRATE USING A SINGLE, DAILY DOSE OF A GAMMA-HYDROXYBUTYRATE FORMULATION
- U-3602 TREATMENT OF EXCESSIVE DAYTIME SLEEPINESS WITH A DOSING REGIMEN THAT INCLUDES ORAL ADMINISTRATION OF 75 MG ONCE DAILY FOR AT LEAST 3 DAYS FOLLOWED BY 150 MG ONCE DAILY
- U-3603 TREATMENT OF CF IN A PATIENT AGE 1 MONTH TO <4 MONTHS WHO HAS AT LEAST ONE MUTATION IN THE CFTR GENE THAT IS RESPONSIVE TO IVACAFTOR BASED ON CLINICAL AND/OR IN VITRO ASSAY DATA USING THE COMPOSITION RECITED IN CLAIM 1 OF US 11564916
- U-3604 TREATMENT OF CF IN A PATIENT AGE 1 MONTH TO <4 MONTHS WHO HAS AT LEAST ONE MUTATION IN THE CFTR GENE THAT IS RESPONSIVE TO IVACAFTOR BASED ON CLINICAL AND/OR IN VITRO ASSAY DATA USING THE COMPOSITION RECITED IN CLAIM 1 OF US 11147770
- U-3605 TREATMENT OF CF IN A PATIENT AGE 1 MONTH TO <4 MONTHS WHO HAS AT LEAST ONE MUTATION IN THE CFTR GENE THAT IS RESPONSIVE TO IVACAFTOR BASED ON CLINICAL AND/OR IN VITRO ASSAY DATA USING THE COMPOSITION RECITED IN CLAIM 1 OF US 10272046
- U-3606 TREATMENT OF CF IN A PATIENT AGE 1 MONTH TO <4 MONTHS WHO HAS AT LEAST ONE CFTR MUTATION RESPONSIVE TO IVACAFTOR BASED ON CLINICAL AND/OR IN VITRO ASSAY DATA USING A SOLID COMPOSITION COMPRISING AMORPHOUS (LESS THAN ABOUT 30% CRYSTALLINE) IVACAFTOR
- U-3607 TREATMENT OF A MODERATE TO MILD CLINICAL PHENOTYPE OF CF USING IVACAFTOR IN A PATIENT AGE 1 MONTH TO <4 MONTHS WHO HAS AT LEAST ONE CFTR MUTATION RESPONSIVE TO IVACAFTOR BASED ON CLINICAL AND/OR IN VITRO ASSAY DATA
- U-3608 TREATMENT OF CYSTIC FIBROSIS USING IVACAFTOR IN A PATIENT AGE 1 MONTH TO <4 MONTHS WHO HAS A R117H MUTATION IN THE CFTR GENE
- U-3609 TREATMENT OF CYSTIC FIBROSIS USING IVACAFTOR IN A PATIENT AGE 1 MONTH TO <4 MONTHS WHO HAS AT LEAST ONE MUTATION IN THE CFTR GENE THAT IS RESPONSIVE TO IVACAFTOR BASED ON CLINICAL AND/OR IN VITRO ASSAY DATA
- U-3610 CO-ADMINISTRATION OF CENOBAMATE WITH PHENOBARBITAL AND/OR PHENYTOIN FOR THE TREATMENT OF PARTIAL ONSET SEIZURES
- U-3611 TREATMENT OF PEDIATRIC PATIENTS WITH CENTRAL PRECOCIOUS PUBERTY
- U-3612 AS AN ADJUNCT TO DIET TO REDUCE LOW-DENSITY LIPOPROTEIN CHOLESTEROL IN ADULTS AND PEDIATRIC PATIENTS AGED 10 YEARS AND OLDER WITH HETEROZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA
- U-3613 AS AN ADJUNCT TO OTHER LOW-DENSITY LIPOPROTEIN CHOLESTEROL (LDL-C) LOWERING THERAPIES, OR ALONE IF SUCH TREATMENTS ARE UNAVAILABLE, TO REDUCE LDL-C IN ADULTS AND PEDIATRIC PATIENTS AGED 10 YEARS AND OLDER WITH HOMOZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA
- U-3614 POSITRON EMISSION TOMOGRAPHY (PET) OF PROSTATE-SPECIFIC MEMBRANE ANTIGEN (PSMA) POSITIVE LESIONS IN MEN WITH PROSTATE CANCER WITH SUSPECTED METASTASIS WHO ARE CANDIDATES FOR INITIAL DEFINITIVE THERAPY
- U-3615 POSITRON EMISSION TOMOGRAPHY (PET) OF PROSTATE-SPECIFIC MEMBRANE ANTIGEN (PSMA) POSITIVE LESIONS IN MEN WITH PROSTATE CANCER WITH SUSPECTED RECURRENCE BASED ON ELEVATED SERUM PROSTATE-SPECIFIC ANTIGEN (PSA) LEVEL
- U-3616 A METHOD OF ADMINISTERING AN OPIOID MAINTENANCE TREATMENT COMPRISING BUPRENORPHINE
- U-3617 A METHOD OF ADMINISTERING AN OPIOID MAINTENANCE TREATMENT COMPRISING BUPRENORPHINE. A METHOD OF TREATING OPIOID WITHDRAWAL USING AN OPIOID

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

MAINTENANCE TREATMENT COMPRISING BUPRENORPHINE

- U-3618 A METHOD OF SUSTAINED DELIVERY OF BUPRENORPHINE TO A HUMAN OR NON-HUMAN ANIMAL BODY. A METHOD FOR TREATMENT FOR OPIOID MAINTENANCE THERAPY
- U-3619 A METHOD OF DELIVERY OF A BIOACTIVE AGENT BY SUBCUTANEOUS INJECTION. A METHOD OF TREATMENT OF A HUMAN FOR ADDICTION
- U-3620 A METHOD OF DELIVERY OF AN OPIOID BIOACTIVE AGENT. A METHOD OF TREATMENT OR PROPHYLAXIS OF A HUMAN OR NON-HUMAN ANIMAL FOR THE TREATMENT OF OPIOID ADDICTION AND/OR THE SYMPTOMS OF OPIOID WITHDRAWAL
- U-3621 TREATMENT OF MODERATE TO SEVERE VASOMOTOR SYMPTOMS DUE WITH MENOPAUSE
- U-3622 TREATMENT OF MODERATE TO SEVERE VASOMETER SYMPTOMS DUE TO MENOPAUSE
- U-3623 MAINTENANCE TREATMENT OF ASTHMA IN PATIENTS AGED 5 YEARS AND OLDER. RECOMMENDED DOSAGES: BREO 100/25 OR 200/25 AGES 18 YEARS AND OLDER; BREO 100/25 AGES 12-17 YEARS, AND BREO 50/25, AGES 5-11 YEARS
- U-3624 TREATMENT OF ADULTS WITH MODERATELY TO SEVERELY ACTIVE CROHN'S DISEASE WHO HAVE HAD AN INADEQUATE RESPONSE OR INTOLERANCE TO ONE OR MORE TNF BLOCKERS
- U-3626 TREATMENT OF SCHIZOPHRENIA BY ADMINISTRATION OF A SIX-MONTH PALIPERIDONE PALMITATE INJECTABLE SUSPENSION FILLED SYRINGE THAT HAS BEEN SHIPPED AND STORED IN A HORIZONTAL POSITION
- U-3627 TREATMENT OF DRY EYE DISEASE (DED)
- U-3628 REDUCTION OF RISK OF CARDIOVASCULAR DEATH, HOSPITALIZATION FOR HEART FAILURE, AND URGENT HEART FAILURE IN ADULTS WITH HEART FAILURE OR TYPE 2 DIABETES MELLITUS, CHRONIC KIDNEY DISEASE, AND OTHER CARDIOVASCULAR RISK FACTORS
- U-3629 TREATMENT OF MILD-TO-MODERATE CORONAVIRUS DISEASE 2019 (COVID-19) IN ADULTS WHO ARE AT HIGH RISK FOR PROGRESSION TO SEVERE COVID-19, INCLUDING HOSPITALIZATION OR DEATH
- U-3630 TREATING OPIOID OVERDOSE
- U-3631 TREATMENT OF DELETERIOUS OR SUSPECTED DELETERIOUS BRCA-MUTATED METASTATIC CASTRATION-RESISTANT PROSTATE CANCER IN COMBINATION WITH ABIRATERONE AND PREDNISONE OR PREDNISOLONE
- U-3632 TREATMENT OF ADULT PATIENTS WITH MANTLE CELL LYMPHOMA
- U-3633 TREATMENT OF ADULT PATIENTS WITH MULTIPLE MYELOMA
- U-3634 METHOD OF TREATING IDA IN ADULT PATIENTS WEIGHING AT LEAST 46.7 KG WITH HEART FAILURE & NY HEART ASSOCIATION CLASS II/III TO IMPROVE EXERCISE CAPACITY BY ADMINISTERING IV FERRIC CARBOXYMALTOSE TO PROVIDE AT LEAST 0.7 G OF ELEMENTAL IRON IN 15 MINUTES
- U-3635 METHOD TO TREAT IDA IN ADULTS WEIGHING AT LEAST 40 KG WITH HEART FAILURE & NY HEART ASSOCIATION CLASS II/III TO IMPROVE EXERCISE CAPACITY BY ADMINISTERING IV FERRIC CARBOXYMALTOSE TO PROVIDE AT LEAST ABOUT 0.6 G OF ELEMENTAL IRON IN ABOUT ≤ 15 MIN
- U-3636 METHOD TO TREAT IDA IN ADULTS WEIGHING AT LEAST 40 KG WITH HEART FAILURE & NY HEART ASSOCIATION CLASS II/III TO IMPROVE EXERCISE CAPACITY BY INTRAVENOUSLY ADMINISTERING FERRIC CARBOXYMALTOSE TO PROVIDE AT LEAST ABOUT 0.6 G OF ELEMENTAL IRON
- U-3637 METHOD TO TREAT IRON DEFICIENCY IN ADULTS WEIGHING AT LEAST 40 KG WITH HEART FAILURE & NY HEART ASSOCIATION CLASS II/III TO IMPROVE EXERCISE CAPACITY BY ADMINISTERING IV FERRIC CARBOXYMALTOSE TO PROVIDE AT LEAST ABOUT 0.6 G OF ELEMENTAL IRON
- U-3638 A METHOD FOR TREATING AND/OR REDUCING THE RISK OF ACUTE MYOCARDIAL INFARCTION
- U-3639 A METHOD FOR TREATING AND/OR REDUCING THE RISK OF A CARDIOVASCULAR EVENT
- U-3640 A METHOD OF TREATING AND/OR REDUCING THE RISK OF INFLAMMATION, ATHEROSCLEROTIC VASCULAR DISEASE, AND CHOLESTEROL CRYSTAL INDUCED INFLAMMATION WITHIN ATHEROSCLEROTIC PLAQUES
- U-3641 A METHOD OF TREATING AND/OR REDUCING THE RISK OF A CARDIOVASCULAR EVENT; ACUTE CORONARY SYNDROME, OUT-OF-HOSPITAL CARDIAC ARREST, AND/OR NONCARDIOEMBOLIC ISCHEMIC STROKE
- U-3642 A METHOD OF TREATING CARDIOVASCULAR DISEASE
- U-3643 METHOD OF TREATING AND/OR REDUCING THE RISK OF A CARDIOVASCULAR EVENT

PATENT AND EXCLUSIVITY TERMS

ADB 206 of 225

PATENT USE

- U-3644 TREATMENT OF FUNCTIONAL CONSTIPATION IN PEDIATRIC PATIENTS 6 TO 17 YEARS OF AGE
- U-3645 ACUTE TREATMENT OF AGITATION ASSOCIATED WITH SCHIZOPHRENIA OR BIPOLAR I OR II DISORDER BY BUCCAL ADMINISTRATION
- U-3646 A METHOD OF TREATMENT OF OVARIAN CANCER OR FALLOPIAN TUBE CANCER
- U-3647 A METHOD OF TREATMENT OF RECURRENT OVARIAN CANCER OR FALLOPIAN TUBE CANCER ASSOCIATED WITH DELETERIOUS OR SUSPECTED DELETERIOUS GERMLINE BRCA-MUTATION
- U-3648 METHOD OF TREATING CHOLESTATIC PRURITUS IN PATIENTS 12 MONTHS OR OLDER SUFFERING FROM ALAGILLE SYNDROME (ALGS)
- U-3649 METHOD OF REDUCING SERUM BILE ACIDS IN PATIENTS 12 MONTHS OR OLDER SUFFERING FROM ALAGILLE SYNDROME (ALGS)
- U-3650 TREATMENT OF PATIENTS WITH POST-TRANSPLANT CYTOMEGALOVIRUS (CMV) INFECTION/DISEASE REFRACTORY TO TREATMENT WITH GANCICLOVIR, VALGANCICLOVIR, CIDOFOVIR, OR FOSCARNET, WHERE THE PATIENT IS A STEM CELL, KIDNEY, OR LIVER TRANSPLANT RECIPIENT
- U-3651 TREATMENT OF ADULT PATIENTS WITH HRR GENE-MUTATED METASTATIC CASTRATION-RESISTANT PROSTATE CANCER (MCRPC) IN COMBINATION WITH ENZALUTAMIDE
- U-3652 AS AN ADJUNCT TO DIET AND STATIN THERAPY FOR THE TREATMENT OF ADULTS WITH PRIMARY HYPERLIPIDEMIA, INCLUDING HETEROZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA (HEFH), BY INHIBITING EXPRESSION OF THE PCSK9 GENE
- U-3653 MANAGEMENT OF MODERATE TO SEVERE PAIN ASSOCIATED WITH ENDOMETRIOSIS IN A PATIENT WITH MODERATE HEPATIC IMPAIRMENT
- U-3654 MANAGEMENT OF MODERATE TO SEVERE PAIN ASSOCIATED WITH ENDOMETRIOSIS USING 150 MG OR 200 MG ELAGOLIX WHILE CO-ADMINISTERING OMEPRAZOLE
- U-3655 MANAGEMENT OF HEAVY MENSTRUAL BLEEDING ASSOCIATED WITH UTERINE LEIOMYOMAS (FIBROIDS) USING 300 MG ELAGOLIX WHILE CO-ADMINISTERING OMEPRAZOLE
- U-3656 TREATMENT OF METASTATIC COLORECTAL CANCER ALONE OR WITH BEVACIZUMAB IN PATIENTS PREVIOUSLY TREATED WITH FLUOROPYRIMIDINE-, OXALIPLATIN- AND IRINOTECAN-BASED CHEMOTHERAPY, AN ANTI-VEGF BIOLOGICAL THERAPY, AND IF RAS WILD-TYPE, AN ANTI-EGFR THERAPY
- U-3657 TREATMENT OF METASTATIC COLORECTAL CANCER ALONE OR WITH BEVACIZUMAB IN SEVERELY RENALLY IMPAIRED PATIENTS PREVIOUSLY TREATED WITH FLUOROPYRIMIDINE-, OXALIPLATIN- AND IRINOTECAN THERAPY, AN ANTI-VEGF BIOLOGIC, AND IF RAS WILD-TYPE, ANTI-EGFR THERAPY
- U-3658 TREATMENT OF METASTATIC GASTRIC OR GJA IN SEVERELY RENALLY IMPAIRED PATIENTS TREATED WITH AT LEAST TWO LINES OF CHEMOTHERAPY THAT INCLUDED A FLUOROPYRIMIDINE, A PLATINUM, A TAXANE OR IRINOTECAN, AND IF APPROPRIATE, HER2/NEU-TARGETED THERAPY
- U-3659 TREATMENT OF METASTATIC COLORECTAL CANCER WITH BEVACIZUMAB IN PATIENTS PREVIOUSLY TREATED WITH FLUOROPYRIMIDINE-, OXALIPLATIN- AND IRINOTECAN-BASED CHEMOTHERAPY, AN ANTI-VEGF BIOLOGICAL THERAPY, AND IF RAS WILD-TYPE, AN ANTI-EGFR THERAPY
- U-3660 TREATMENT OF PARTIAL-ONSET SEIZURES IN ADULTS AND IN PEDIATRIC PATIENTS WEIGHING AT LEAST 50 KG
- U-3661 COMBINATION WITH STANDARD CYTARABINE AND ANTHRACYCLINE INDUCTION AND CYTARABINE CONSOLIDATION, AND AS MAINTENANCE MONOTHERAPY FOLLOWING CONSOLIDATION CHEMOTHERAPY, FOR ADULT PATIENTS WITH NEWLY DIAGNOSED FLT3-ITD POSITIVE ACUTE MYELOID LEUKEMIA
- U-3662 DEXTROMETHORPHAN AND BUPROPRION IN COMBINATION TO TREAT MAJOR DEPRESSIVE DISORDER
- U-3663 TOPICAL TREATMENT OF MOLLUSCUM CONTAGIOSUM IN ADULT AND PEDIATRIC PATIENTS 2 YEARS OF AGE AND OLDER
- U-3664 TOPICAL TREATMENT OF SKIN LESIONS CAUSED BY AN INFECTION WITH MOLLUSCUM CONTAGIOSUM IN ADULT AND PEDIATRIC PATIENTS 2 YEARS OF AGE AND OLDER
- U-3665 TOPICAL DELIVERY OF A CANTHARIDIN FORMULATION TO ADULT AND PEDIATRIC PATIENTS 2 YEARS OF AGE AND OLDER FOR TREATING MOLLUSCUM CONTAGIOSUM
- U-3666 USE IN ADULT AND PEDIATRIC PATIENTS WITH SUBOPTIMAL ECHOCARDIOGRAMS TO OPACIFY THE LEFT VENTRICULAR CHAMBER AND TO IMPROVE THE DELINEATION OF THE LEFT VENTRICULAR ENDOCARDIAL BORDER
- U-3667 USE IN ULTRASONOGRAPHY OF THE URINARY TRACT IN PEDIATRIC PATIENTS FOR THE

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

- EVALUATION OF SUSPECTED OR KNOWN VESICoureTERAL REFLUX
- U-3668 USE WITH ULTRASOUND OF THE LIVER IN ADULT AND PEDIATRIC PATIENTS TO CHARACTERIZE FOCAL LIVER LESIONS
- U-3669 USE OF VASCEPA TO REDUCE THE INCIDENCE OF STROKE IN AN ADULT PATIENT ON STATIN THERAPY AND WITH ELEVATED TRIGLYCERIDE LEVELS (>150 MG/DL), WHEREIN THE PATIENT EXPERIENCES ATRIAL FIBRILLATION AND/OR FLUTTER INSTEAD OF AN INCIDENCE OF STROKE
- U-3670 ADMINISTRATION OF AN EXTENDED RELEASE TABLET FOR THE TREATMENT OF NEUROGENIC DETRUSOR OVERACTIVITY (NDO) IN PEDIATRIC PATIENTS AGED 3 YEARS AND OLDER AND WEIGHING 35 KG OR MORE
- U-3671 USE OF INTRANASAL NALOXONE FOR THE TREATMENT OF OPIOID OVERDOSE
- U-3672 MANAGEMENT OF MODERATE TO SEVERE PAIN ASSOCIATED WITH ENDOMETRIOSIS USING 150 MG ELAGOLIX FOR UP TO 24 MONTHS
- U-3673 TREATMENT OF GEOGRAPHIC ATROPHY SECONDARY TO AGE-RELATED MACULAR DEGENERATION BY ADMINISTERING AVACINCAPTAD PEGOL TO THE EYE
- U-3674 TREATMENT OF DEMODEX BLEPHARITIS VIA TOPICAL ADMINISTRATION TO AN OCULAR SURFACE
- U-3675 A PERCUTANEOUS HEPATIC PERFUSION PROCEDURE FOR TREATING A PATIENT WITH UVEAL MELANOMA WITH UNRESECTABLE HEPATIC METASTASES
- U-3676 REDUCTION OF HETEROTOPIC OSSIFICATION IN PATIENTS WITH FIBRODYSPLASIA (MYOSITIS) OSSIFICANS PROGRESSIVA
- U-3677 ACUTE TREATMENT OF MIGRAINE WITH OR WITHOUT AURA IN A PATIENT WITH SEVERE HEPATIC IMPAIRMENT
- U-3678 A METHOD FOR REDUCING THE INCIDENCE OF CHEMOTHERAPY-INDUCED MYELOSUPPRESSION IN ADULT PATIENTS WHEN ADMINISTERED PRIOR TO A PLATINUM/ETOPOSIDE-CONTAINING REGIMEN FOR EXTENSIVE-STAGE SMALL CELL LUNG CANCER
- U-3679 A METHOD FOR REDUCING THE INCIDENCE OF CHEMOTHERAPY-INDUCED MYELOSUPPRESSION IN ADULT PATIENTS WHEN ADMINISTERED PRIOR TO A TOPOTECAN-CONTAINING REGIMEN FOR EXTENSIVE-STAGE SMALL CELL LUNG CANCER
- U-3680 A METHOD FOR TREATING A SUBJECT WITH UVEAL MELANOMA WITH UNRESECTABLE HEPATIC METASTASES
- U-3681 TREATMENT OF ACUTE CYANIDE POISONING THAT IS JUDGED TO BE SERIOUS OR LIFE-THREATENING
- U-3682 SODIUM THIOSULFATE INJECTION IS INDICATED FOR SEQUENTIAL USE WITH SODIUM NITRITE FOR THE TREATMENT OF ACUTE CYANIDE POISONING THAT IS JUDGED TO BE SERIOUS OR LIFE-THREATENING
- U-3683 A METHOD OF TREATING A PATIENT WITH UVEAL MELANOMA WITH UNRESECTABLE HEPATIC METASTASES
- U-3684 A METHOD OF TREATING AN ADULT PATIENT WITH RELAPSED OR REFRACTORY ACUTE MYELOID LEUKEMIA HAVING A SUSCEPTIBLE IDH1 MUTATION
- U-3685 METHOD OF ADMINISTERING AN EFFECTIVE DOSE OF TROPICAMIDE AND PHENYLEPHRINE HYDROCHLORIDE TO AN EYE
- U-3686 TREATMENT OF EXACERBATIONS OF MULTIPLE SCLEROSIS IN ADULTS WITH CORTICOTROPIN BY PROMOTING NEW VESSEL FORMATION WHEREIN VCAM-1 EXPRESSION AND ANGIOPOETIN-2 EXPRESSION IS INCREASED AFTER THE ADMINISTERING
- U-3687 TREATMENT OF INFANTILE SPASMS WITH CORTICOTROPIN BY PROMOTING NEW VESSEL FORMATION WHEREIN VCAM-1 EXPRESSION AND ANGIOPOETIN-2 EXPRESSION IS INCREASED AFTER THE ADMINISTERING
- U-3688 TREATMENT OF OPHTHALMIC DISEASES WITH CORTICOTROPIN BY PROMOTING NEW VESSEL FORMATION WHEREIN VCAM-1 EXPRESSION AND ANGIOPOETIN-2 EXPRESSION IS INCREASED AFTER THE ADMINISTERING
- U-3689 DIAGNOSIS OF GALL BLADDER DISORDERS BY STIMULATING GALLBLADDER CONTRACTION, AS MAY BE ASSESSED BY VARIOUS METHODS OF DIAGNOSTIC IMAGING, OR TO OBTAIN BY DUODENAL ASPIRATION A SAMPLE OF CONCENTRATED BILE FOR ANALYSIS
- U-3690 A METHOD FOR PREVENTING OF POST-OPERATIVE NAUSEA AND VOMITING
- U-3691 TREATMENT AND REDUCTION OF RISK BY ADMINISTRATION OF EMPAGLIFLOZIN TO ADULTS WITH CHRONIC KIDNEY DISEASE AT RISK OF PROGRESSION
- U-3692 A METHOD OF TREATING HETEROZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA OR ESTABLISHED ATHEROSCLEROTIC CARDIOVASCULAR DISEASE BY DECREASING THE LEVEL OF LDL-C USING 180 MG BEMPEDOIC ACID AND 10 MG EZETIMIBE

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

- U-3693 TREATMENT OF EXCESSIVE DAYTIME SLEEPINESS IN A BREAST-FEEDING PATIENT WHILE REDUCING INFANT EXPOSURE TO SOLRIAMFETOL
- U-3694 METHOD OF REDUCING OR AMELIORATING SEIZURES IN A PATIENT BY ADMINISTERING A LIQUID FORMULATION OF FENFLURAMINE WITH STIRIPENTOL THEREBY MODULATING DOWN THE FORMATION OF NORFENFLURAMINE AND RESULTING IN HIGHER LEVELS OF FENFLURAMINE
- U-3695 MAINTENANCE TREATMENT OF DELETERIOUS OR SUSPECTED DELETERIOUS GERMLINE OR SOMATIC BRCA-MUTED RECURRENT EPITHELIAL OVARIAN, FALLOPIAN TUBE OR PRIMARY PERITONEAL CANCER, WHO ARE IN A COMPLETE OR PARTIAL RESPONSE TO PLATINUM-BASED CHEMOTHERAPY
- U-3696 TREATMENT OF CF IN PATIENTS AGED 2 TO <6 YEARS WHO HAVE AT LEAST ONE F508DEL MUTATION OR A MUTATION IN THE CFTR GENE THAT IS RESPONSIVE BASED ON IN VITRO DATA COMPRISING ADMINISTERING A COMPOSITION ACCORDING TO CLAIM 1 OF US 11752106
- U-3697 TREATMENT OF CF IN A PATIENT AGE 1 MONTH TO <6 YEARS WHO HAS AT LEAST ONE MUTATION IN THE CFTR GENE THAT IS RESPONSIVE TO IVACAFTOR BASED ON CLINICAL AND/OR IN VITRO ASSAY DATA USING THE COMPOSITION RECITED IN CLAIM 1 OF US 11752106
- U-3698 ACUTE TREATMENT OF AGITATION ASSOCIATED WITH SCHIZOPHRENIA BY SUBLINGUAL OR BUCCAL ADMINISTRATION
- U-3699 TREATMENT OF SUBJECTS WITH MAJOR DEPRESSION WITH SEXUAL DYSFUNCTION CAUSED BY EITHER MAJOR DEPRESSION OR PRIOR TREATMENTS, OR TREATMENT OF SUBJECTS WITH MAJOR DEPRESSION WITHOUT THE RISK OF SEXUAL DYSFUNCTION ADVERSE REACTIONS
- U-3700 TREATMENT OF ADULTS WITH ACUTE LYMPHOBLASTIC LEUKEMIA WITH AN ORAL SOLUTION OF METHOTREXATE
- U-3701 TREATMENT OF ADULTS WITH MYCOSIS FUNGOIDES WITH AN ORAL SOLUTION OF METHOTREXATE
- U-3702 TREATMENT OF ADULTS WITH RELAPSED OR REFRACTORY NON-HODGKIN LYMPHOMAS WITH AN ORAL SOLUTION OF METHOTREXATE
- U-3703 TREATMENT OF ADULTS WITH RHEUMATOID ARTHRITIS WITH AN ORAL SOLUTION OF METHOTREXATE
- U-3704 TREATMENT OF ADULTS WITH SEVERE PSORIASIS WITH AN ORAL SOLUTION OF METHOTREXATE
- U-3705 TREATMENT OF CATAPLEXY OR EXCESSIVE DAYTIME SLEEPINESS IN PATIENTS WITH NARCOLEPSY WHO HAVE AN APNEA/HYPOPNEA INDEX ≤ 15 WITH A ONCE-NIGHTLY FORMULATION OF GAMMA-HYDROXYBUTYRATE
- U-3706 TREATMENT OF BLOATING ASSOCIATED WITH DIARRHEA-PREDOMINANT IRRITABLE BOWEL SYNDROME (IBS-D) IN ADULT FEMALE SUBJECTS
- U-3707 A METHOD OF TREATING PATIENTS 1 YEAR OF AGE AND OLDER WITH CHRONIC PHASE PH+ CML, NEWLY-DIAGNOSED OR RESISTANT OR INTOLERANT TO PRIOR THERAPY
- U-3708 A METHOD OF TREATING PATIENTS WITH ACCELERATED, OR BLAST PHASE PH+ CML WITH RESISTANCE OR INTOLERANCE TO PRIOR THERAPY
- U-3709 METHOD OF TREATING PRIMARY HYPEROXALURIA TYPE 1 (PH1)
- U-3710 TREATMENT OF ADULT PATIENTS WITH PREVIOUSLY UNTREATED CHRONIC LYMPHOCYTIC LEUKEMIA BY ORALLY ADMINISTERING 100 MG OF ACALABRUTINIB TWICE DAILY IN COMBINATION WITH INTRAVENOUS ADMINISTRATION OF OBINUTUZUMAB
- U-3711 A METHOD TO IMPROVE KIDNEY FUNCTION IN ADULTS WITH HEPATORENAL SYNDROME WITH RAPID REDUCTION IN KIDNEY FUNCTION
- U-3712 TOPICAL TREATMENT OF PLAQUE PSORIASIS, INCLUDING INTERTRIGINOUS AREAS, IN PATIENTS 6 YEARS OF AGE OR OLDER
- U-3713 TOPICAL TREATMENT OF ACNE VULGARIS IN ADULT AND PEDIATRIC PATIENTS 12 YEARS OF AGE AND OLDER
- U-3714 TREATMENT OF GASTROINTESTINAL STROMAL TUMORS IN PATIENTS HAVING PRE-EXISTING SEVERE HEPATIC IMPAIRMENT AND SUFFERING FROM AN ADVERSE EVENT WHILE BEING ADMINISTERED RIPRETINIB DAILY
- U-3715 TREATMENT OF ADULT PATIENTS WITH CHRONIC LYMPHOCYTIC LEUKEMIA (CLL) OR SMALL LYMPHOCYTIC LYMPHOMA (SLL) RECEIVING A MODERATE CYP3A INDUCER
- U-3716 TREATMENT OF ADULT PATIENTS WITH WALDENSTROM'S MACROGLOBULINEMIA (WM) RECEIVING A MODERATE CYP3A INDUCER
- U-3717 TREATMENT OF ADULT PATIENTS WITH SMALL LYMPHOCYTIC LYMPHOMA (SLL) RECEIVING A MODERATE CYP3A INDUCER
- U-3718 TREATMENT OF ADULT PATIENTS WITH RELAPSED OR REFRACTORY MARGINAL ZONE LYMPHOMA

PATENT AND EXCLUSIVITY TERMS

ADB 209 of 225

PATENT USE

- (MZL) RECEIVING A MODERATE CYP3A INDUCER, WHO HAVE RECEIVED AT LEAST ONE ANTI-CD20-BASED REGIMEN
- U-3719 TREATMENT OF ADULT PATIENTS WITH MANTLE CELL LYMPHOMA (MCL) RECEIVING A MODERATE CYP3A INDUCER, WHO HAVE RECEIVED AT LEAST ONE PRIOR THERAPY
- U-3720 TREATMENT OF ADULT PATIENTS WITH CHRONIC LYMPHOCYTIC LEUKEMIA (CLL) RECEIVING A MODERATE CYP3A INDUCER
- U-3721 TREATMENT OF SECONDARY HYPERPARATHYROIDISM WITH SUSTAINED RELEASE 25-HYDROXYVITAMIN D IN CHRONIC KIDNEY DISEASE PATIENTS RECEIVING CHOLESTYRAMINE
- U-3722 TREATMENT OF SECONDARY HYPERPARATHYROIDISM WITH SUSTAINED RELEASE 25-HYDROXYVITAMIN D IN CHRONIC KIDNEY DISEASE PATIENTS RECEIVING PHENOBARBITAL OR OTHER ANTICONVULSANTS
- U-3723 TREATMENT OF SECONDARY HYPERPARATHYROIDISM WITH SUSTAINED RELEASE 25-HYDROXYVITAMIN D IN CHRONIC KIDNEY DISEASE PATIENTS RECEIVING CYP3A INHIBITORS
- U-3724 TO PRODUCE POST-SURGICAL ANALGESIA
- U-3725 ACUTE TREATMENT OF AGITATION ASSOCIATED WITH SCHIZOPHRENIA OR BIPOLAR I OR II DISORDER BY SUBLINGUAL OR BUCCAL ADMINISTRATION IN PATIENTS WITH SEVERE HEPATIC IMPAIRMENT
- U-3726 THE TREATMENT OF POMPE PATIENTS
- U-3727 TREATMENT OF ADULT PATIENTS WITH SMALL LYMPHOCYTIC LYMPHOMA (SLL) WHO HAVE RECEIVED AT LEAST ONE PRIOR THERAPY
- U-3728 TREATMENT OF ADULT PATIENTS WITH CHRONIC LYMPHOCYTIC LEUKEMIA (CLL) WHO HAVE RECEIVED AT LEAST ONE PRIOR THERAPY
- U-3729 TREATMENT OF ADULT PATIENTS WITH CHRONIC LYMPHOCYTIC LEUKEMIA (CLL) OR SMALL LYMPHOCYTIC LYMPHOMA (SLL) WHO HAVE RECEIVED AT LEAST ONE PRIOR THERAPY
- U-3730 A METHOD FOR TREATING ULCERATIVE COLITIS BY ADMINISTERING ESTRASIMOD ARGININE IN AN AMOUNT EQUIVALENT TO ABOUT 2.0 MG OF ESTRASIMOD
- U-3731 A METHOD FOR TREATING ULCERATIVE COLITIS BY ADMINISTERING A THERAPEUTICALLY EFFECTIVE AMOUNT OF THE FORM OF ESTRASIMOD ARGININE AS CLAIMED
- U-3732 A METHOD FOR TREATING ULCERATIVE COLITIS BY ADMINISTERING A THERAPEUTICALLY EFFECTIVE AMOUNT OF ESTRASIMOD ARGININE
- U-3733 METHOD OF TREATING SEIZURES IN A PATIENT BY ADMINISTERING A LIQUID FORMULATION OF FENFLURAMINE OR ITS SALTS PLUS STIRIPENTOL THEREBY REDUCING NORFENFLURAMINE FORMATION WHILE INCREASING THE FENFLURAMINE LEVEL. PATIENTS CAN HAVE E.G. DRAVET OR LGS
- U-3734 METHOD OF TREATING SCHIZOPHRENIA IN A PATIENT WHO HAS PREVIOUSLY EXPERIENCED SIGNIFICANT WEIGHT GAIN INDUCED BY OLANZAPINE ALONE BY ADMINISTERING A COMPOSITION COMPRISING OLANZAPINE AND SAMIDORPHAN
- U-3735 TREATMENT OF GENERALIZED MYASTHENIA GRAVIS (GMG) IN AN ADULT PATIENT WHO IS ANTI-ACETYLCHOLINE RECEPTOR (ACHR) ANTIBODY POSITIVE BY SUBCUTANEOUS ADMINISTRATION OF C5 COMPLEMENT INHIBITOR ZILUCOPLAN
- U-3736 REDUCTION OF SERUM PHOSPHORUS IN ADULTS
- U-3737 MEKTOVI IS A KINASE INHIBITOR INDICATED IN COMBINATION WITH ENCORAFENIB, FOR THE TREATMENT OF PATIENTS WITH METASTATIC NON SMALL CELL LUNG CANCER WITH A BRAF V600E MUTATION, AS DETECTED BY AN FDA-APPROVED TEST
- U-3738 BRAFTOVI IS A KINASE INHIBITOR INDICATED IN COMBINATION WITH BINIMETINIB, FOR THE TREATMENT OF PATIENTS WITH METASTATIC NON SMALL CELL LUNG CANCER WITH A BRAF V600E MUTATION, AS DETECTED BY AN FDA-APPROVED TEST
- U-3739 TREATMENT OF NON-24-HOUR SLEEP-WAKE DISORDER BY AVOIDING THE ADMINISTRATION OF TASIMELTEON WITH FOOD
- U-3740 TREATMENT OF MODERATELY TO SEVERELY ACTIVE ULCERATIVE COLITIS (UC) IN ADULTS
- U-3741 TREATMENT OF PRESBYOPIA
- U-3742 A METHOD OF TREATING A CANCER CHARACTERIZED BY AN IDH1 MUTATION WHEREIN THE CANCER IS RELAPSED OR REFRACTORY MYELODYSPLASTIC SYNDROMES
- U-3743 A METHOD OF TREATING RELAPSED OR REFRACTORY MYELODYSPLASTIC SYNDROMES CHARACTERIZED BY THE PRESENCE OF A MUTANT ALLELE OF IDH1 BY ADMINISTERING A ONCE DAILY 500 MG ORAL DOSE TO A SUBJECT THAT HAS NOT INGESTED A HIGH-FAT MEAL
- U-3744 METHOD OF TREATING MILD TO MODERATE PAIN IN ADULTS

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

- U-3745 METHOD OF TREATING MODERATE TO SEVERE PAIN IN ADULTS AS AN ADJUNCT TO OPIOID ANALGESICS
- U-3746 METHODS OF MAKING AQUEOUS COMPOSITION AND TREATING PAIN, INFLAMMATION, FEVER, PATENT DUCTUS ATERIOSIS WITH AQUEOUS COMPOSITION
- U-3747 TREATMENT OF DUCHENNE MUSCULAR DYSTROPHY
- U-3748 TOPICAL TREATMENT OF PLAQUE PSORIASIS, INCLUDING INTERTRIGINOUS AREAS, IN PATIENTS 6 YEARS OF AGE AND OLDER
- U-3749 METHOD OF TREATING PULMONARY HYPERTENSION ASSOCIATED WITH INTERSTITIAL LUNG DISEASE BY ADMINISTERING TREPROSTINIL OR A SALT THEREOF BY INHALATION USING A DEVICE
- U-3750 TREATMENT OF GASTROINTESTINAL STROMAL TUMOR WITH A WILD TYPE KIT MUTATION IN PATIENTS PREVIOUSLY ADMINISTERED THREE OR MORE KINASE INHIBITORS, WHERE ONE OF THE KINASE INHIBITORS IS IMATINIB
- U-3751 TREATMENT OF CATAPLEXY OR EXCESSIVE DAYTIME SLEEPINESS IN PATIENTS WITH NARCOLEPSY WITH A ONCE-NIGHTLY GAMMA-HYDROXYBUTYRATE FORMULATION
- U-3752 METHOD OF TREATING DIABETIC HYPOGLYCEMIA
- U-3753 TREATMENT OF PATIENTS WITH METASTATIC COLORECTAL CANCER WHO HAVE BEEN PREVIOUSLY TREATED WITH FLUOROPYRIMIDINE-, OXALIPLATIN-, AND IRINOTECAN-BASED CHEMOTHERAPY, ANTI-VEGF THERAPY, AND, IF RAS WILD-TYPE AND MEDICALLY APPROPRIATE, ANTI-EGFR THERAPY
- U-3754 TREATMENT OF ADULT PATIENTS WITH PROGRESSING DESMOID TUMORS
- U-3755 TREATMENT OF ADULT PATIENTS WITH LOCALLY ADVANCED OR METASTATIC ROS1-POSITIVE NON-SMALL CELL LUNG CANCER (NSCLC)
- U-3756 ACUTE TREATMENT OF AGITATION ASSOCIATED WITH BIPOLAR I OR II DISORDER BY SUBLINGUAL OR BUCCAL ADMINISTRATION
- U-3757 TREATMENT OF HYPOTHYROIDISM BY ORAL ADMINISTRATION OF L-THYROXINE TO A PATIENT ON AN EMPTY STOMACH 15 MINUTES BEFORE BREAKFAST
- U-3758 TREATMENT OF PITUITARY THYROTROPIN SUPPRESSION BY ORAL ADMINISTRATION OF L-THYROXINE TO A PATIENT ON AN EMPTY STOMACH 15 MINUTES BEFORE BREAKFAST
- U-3759 METHOD FOR REDUCING THE RISK OF CARDIOVASCULAR DEATH AND HOSPITALIZATION FOR HEART FAILURE IN PATIENTS WITH HEART FAILURE AND TYPE 2 DIABETES MELLITUS BY ADMINISTRATION OF EMPAGLIFLOZIN
- U-3760 METHOD FOR REDUCING THE RISK OF CARDIOVASCULAR DEATH AND HOSPITALIZATION IN PATIENTS WITH HEART FAILURE BY ADMINISTRATION OF EMPAGLIFLOZIN
- U-3761 ADULT PATIENTS WITH CHRONIC LYMPHOCYTIC LEUKEMIA OR SMALL LYMPHOCYTIC LYMPHOMA (CLL/SLL) WHO HAVE RECEIVED AT LEAST TWO PRIOR LINES OF THERAPY, INCLUDING A BTK INHIBITOR AND A BCL-2 INHIBITOR
- U-3762 TREATMENT WITH FULVESTRANT OF HR-POS. HER2-NEG. LOCALLY ADVANCED OR METASTATIC BREAST CANCER WITH PIK3CA/AKT1/PTEN-ALTERATION(S) FOLLOWING PROGRESSION ON ENDOCRINE THERAPY IN THE METASTATIC SETTING OR RECURRENCE ON OR WITHIN 12 MONTHS OF ADJUVANT THERAPY
- U-3763 TREATMENT OF PATIENTS WITH NON-METASTATIC CASTRATION-SENSITIVE PROSTATE CANCER (NMCSPC) WITH BIOCHEMICAL RECURRENCE (BCR) AT HIGH RISK FOR METASTASIS
- U-3764 TREATMENT OF EXCESSIVE DAYTIME SLEEPINESS BY ADMINISTERING SOLRIAMFETOL TO A SUBJECT HAVING OBSTRUCTIVE SLEEP APNEA (OSA) AND NO, MILD, MODERATE, OR SEVERE RENAL IMPAIRMENT
- U-3765 TREATMENT OF EXCESSIVE DAYTIME SLEEPINESS BY ADMINISTERING SOLRIAMFETOL TO A SUBJECT HAVING MODERATE OR SEVERE RENAL IMPAIRMENT
- U-3766 REDUCE THE RISK OF CARDIOVASCULAR DEATH AND HOSPITALIZATION FOR HEART FAILURE AND URGENT HEART FAILURE VISITS IN ADULTS WITH HEART FAILURE WITH PRESERVED EJECTION FRACTION AND WITHOUT TYPE II DIABETES
- U-3767 USE OF DASATINIB FOR TREATMENT OF ADULTS WITH CHRONIC, ACCELERATED, OR MYELOID OR LYMPHOID BLAST PHASE PH+ CML WITH RESISTANCE OR INTOLERANCE TO PRIOR THERAPY INCLUDING IMATINIB, WHEN ADMINISTERED AT ELEVATED GASTRIC PH
- U-3768 USE OF DASATINIB FOR TREATMENT OF NEWLY DIAGNOSED ADULTS WITH PH+ CML IN CHRONIC PHASE, WHEN ADMINISTERED AT ELEVATED GASTRIC PH
- U-3769 USE OF DASATINIB FOR TREATMENT OF ADULTS WITH PH+ ALL WITH RESISTANCE OR INTOLERANCE TO PRIOR THERAPY, WHEN ADMINISTERED AT ELEVATED GASTRIC PH

PATENT AND EXCLUSIVITY TERMS

ADB 211 of 225

PATENT USE

- U-3770 USE OF DASATINIB FOR TREATMENT OF ADULTS WITH CHRONIC, ACCELERATED, OR MYELOID OR LYMPHOID BLAST PHASE PH+ CML WITH RESISTANCE OR INTOLERANCE TO PRIOR THERAPY INCLUDING IMATINIB, WHEN COADMINISTERED WITH A GASTRIC ACID REDUCING AGENT
- U-3771 USE OF DASATINIB FOR TREATMENT OF ADULTS WITH PH+ ALL WITH RESISTANCE OR INTOLERANCE TO PRIOR THERAPY, WHEN COADMINISTERED WITH A GASTRIC ACID REDUCING AGENT
- U-3772 USE OF DASATINIB FOR TREATMENT OF NEWLY DIAGNOSED ADULTS WITH PH+ CML IN CHRONIC PHASE, WHEN COADMINISTERED WITH A GASTRIC ACID REDUCING AGENT
- U-3773 TOPICAL TREATMENT OF SEBORRHEIC DERMATITIS IN PATIENTS 9 YEARS OF AGE AND OLDER
- U-3774 A CATHETER LOCK SOLUTION TO REDUCE CATHETER-RELATED BLOODSTREAM INFECTIONS IN ADULT PATIENTS RECEIVING HEMODIALYSIS THROUGH A CENTRAL VENOUS CATHETER
- U-3775 TREATMENT OF EXCESSIVE DAYTIME SLEEPINESS BY ADMINISTERING SOLRIAMFETOL TO A SUBJECT HAVING NO, MILD, MODERATE, OR SEVERE RENAL IMPAIRMENT
- U-3776 TREATING TYPE 2 DIABETES MELLITUS BY ASSESSING RENAL FUNCTION AND ORALLY ADMINISTERING EMPAGLIFLOZIN IN A DAILY AMOUNT OF 10 MG OR 25 MG IF THE EGFR IS ≥ 30 ML/MIN/1.73 M² AND < 60 ML/MIN/1.73 M², WHEREIN THE TREATMENT IMPROVES GLYCEMIC CONTROL
- U-3777 TREATING TYPE 2 DIABETES MELLITUS BY ASSESSING RENAL FUNCTION AND ORALLY ADMINISTERING EMPAGLIFLOZIN IN A DAILY AMOUNT OF 10 MG OR 25 MG IF THE EGFR ≥ 45 ML/MIN/1.73 M² AND < 60 ML/MIN/1.73 M², WHEREIN THE TREATMENT IMPROVES GLYCEMIC CONTROL
- U-3778 TREATMENT OF MAJOR DEPRESSIVE DISORDER BY ADMINISTERING DEXTROMETHORPHAN AND BUPROPION TO A SUBJECT HAVING MODERATE HEPATIC IMPAIRMENT
- U-3779 METHOD OF TREATING NEONATAL SEIZURES
- U-3780 TREATMENT OF ADULT PATIENTS WITH ADVANCED RENAL CELL CARCINOMA FOLLOWING A PROGRAMMED DEATH RECEPTOR-1 OR PROGRAMMED DEATH-LIGAND INHIBITOR AND A VASCULAR ENDOTHELIAL GROWTH FACTOR TYROSINE KINASE INHIBITOR
- U-3781 REDUCTION IN LOSS OF KIDNEY FUNCTION IN ADULTS WITH PRIMARY IMMUNOGLOBULIN A NEPHROPATHY (IGAN) WHO ARE AT RISK OF DISEASE PROGRESSION
- U-3782 METHOD FOR ADMINISTERING MITAPIVAT OR A SALT OF MITAPIVAT TO MITIGATE DRUG INTERACTIONS IN PATIENTS WITH HEMOLYTIC ANEMIA THAT ARE TAKING MODERATE CYP3A INDUCERS
- U-3783 COMBINATION TREATMENT WITH TUCATINIB AND TRASTUZUMAB OF ADULTS WITH RAS WILD-TYPE HER2-POSITIVE UNRESECTABLE OR METASTATIC COLORECTAL CANCER THAT HAS PROGRESSED FOLLOWING PREVIOUS TREATMENT AS CLAIMED
- U-3784 USE OF ORAL OCTREOTIDE IN COMBINATION WITH A H₂-RECEPTOR ANTAGONIST OR ANTACID FOR LONG-TERM MAINTENANCE TREATMENT IN ACROMEGALY PATIENTS WHO HAVE RESPONDED TO AND TOLERATED TREATMENT WITH OCTREOTIDE OR LANREOTIDE
- U-3785 METHOD OF PREVENTING PREGNANCY BY INSERTING A VAGINAL SYSTEM CONTAINING 103 MG OF SEGESTERONE ACETATE AND 17.4 MG OF ETHINYL ESTRADIOL FOR UP TO THIRTEEN 21/7-DAY (IN/OUT) CYCLES
- U-3786 ACUTE TREATMENT OF MIGRAINE WITH OR WITHOUT AURA IN A PATIENT WITH SEVERE RENAL IMPAIRMENT
- U-3787 A METHOD FOR PREVENTING POST-OPERATIVE NAUSEA AND VOMITING
- U-3788 METHOD OF ADMINISTERING A NITRIC OXIDE RELEASING ACTIVE PHARMACEUTICAL INGREDIENT TO TREAT AND/OR PREVENT VIRAL INFECTION
- U-3789 METHOD OF TREATING AND/OR PREVENTING VIRAL INFECTION USING A NITRIC OXIDE RELEASING ACTIVE INGREDIENT
- U-3790 METHOD OF TREATING AND/OR PREVENTING MOLLUSCUM CONTAGIOSUM WITH A NITRIC OXIDE RELEASING TOPICAL COMPOSITION
- U-3791 METHOD OF TREATING, PREVENTING, OR REDUCING LESIONS CAUSED BY MOLLUSCUM CONTAGIOSUM
- U-3792 METHOD OF TREATING AND/OR PREVENTING VIRAL INFECTION WITH A TOPICAL NITRIC OXIDE RELEASING COMPONENT
- U-3793 METHOD OF ADMINISTERING A NITRIC OXIDE RELEASING API IN A COMBINATION TOPICAL COMPOSITION
- U-3794 METHOD OF TREATING AND/OR PREVENTING VIRAL INFECTION WITH A TOPICAL COMPOSITION INCLUDING A NITRIC OXIDE RELEASING API

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

- U-3795 METHOD OF TREATING AND/OR PREVENTING MOLLUSCUM CONTAGIOSUM WITH A TOPICAL COMPOSITION INCLUDING A NITRIC OXIDE RELEASING API
- U-3796 METHOD OF PREVENTING AND/OR REDUCING APPEARANCE AND/OR SIZE OF MALIGNANT LESION WITH A TOPICAL COMPOSITION INCLUDING A NITRIC OXIDE RELEASING API
- U-3797 METHOD OF TOPICALLY REDUCING LESIONS WITH TWO SEPARATELY STORED COMPONENTS
- U-3798 METHOD OF TOPICALLY REDUCING LESIONS WITH TWO SEPARATELY STORED COMPONENTS WHERE ONE COMPONENT INCLUDES A NITRIC OXIDE RELEASING COMPOUND
- U-3799 METHOD OF TOPICALLY REDUCING LESIONS WITH TWO SEPARATELY STORED COMPONENTS WHERE ONE COMPONENT INCLUDES WATER
- U-3800 METHOD OF APPLYING RELEASED NITRIC OXIDE TO SKIN FROM COMBINATION INCLUDING ANHYDROUS ALCOHOL GEL
- U-3801 METHOD OF INCREASING RELEASE OF NITRIC OXIDE FROM ANHYDROUS ALCOHOL GEL
- U-3802 METHOD OF TREATING SKIN AILMENT WITH NITRIC OXIDE RELEASING MACROMOLECULES AND HYDROPHILIC GEL
- U-3803 METHOD OF APPLICATION OF TOPICAL PHARMACEUTICAL COMPOSITION TO TREAT DERMATOLOGICAL CONDITION
- U-3804 TREATMENT OF PHARMACOLOGICALLY-INDUCED MYDRIASIS
- U-3805 TREATMENT OF ADULTS WITH LOCALLY ADVANCED OR METASTATIC UROTHELIAL CARCINOMA WITH SUSCEPTIBLE FGFR3 GENETIC ALTERATIONS WHOSE DISEASE HAS PROGRESSED ON OR AFTER AT LEAST ONE LINE OF PRIOR SYSTEMIC THERAPY
- U-3806 TREATMENT OF ADULTS WITH METASTATIC UROTHELIAL CARCINOMA WITH SUSCEPTIBLE FGFR3 GENETIC ALTERATIONS WHOSE DISEASE HAS PROGRESSED ON OR AFTER AT LEAST ONE LINE OF PRIOR SYSTEMIC THERAPY
- U-3807 TREATMENT OF ADULTS WITH LOCALLY ADVANCED OR METASTATIC, SURGICALLY UNRESECTABLE UROTHELIAL CARCINOMA WITH SUSCEPTIBLE FGFR3 GENETIC ALTERATIONS WHOSE DISEASE HAS PROGRESSED ON OR AFTER AT LEAST ONE LINE OF PRIOR SYSTEMIC THERAPY
- U-3808 TREATMENT OF ADULTS WITH LOCALLY ADVANCED OR METASTATIC UROTHELIAL CARCINOMA WITH SUSCEPTIBLE FGFR3 GENETIC ALTERATIONS WHOSE DISEASE HAS PROGRESSED ON OR AFTER AT LEAST ONE LINE OF PRIOR SYSTEMIC THERAPY, WITH DOSING BASED ON SERUM PHOSPHATE LEVELS
- U-3809 IN COMBINATION WITH FULVESTRANT FOR TREATMENT OF ADULTS WITH HR-POSITIVE, HER-2-NEGATIVE, PIK3CA-MUTATED, ADVANCED OR METASTATIC BREAST CANCER
- U-3810 REDUCTION IN LOSS OF KIDNEY FUNCTION AND REDUCTION OF PROTEINURIA IN ADULTS WITH PRIMARY IMMUNOGLOBULIN A NEPHROPATHY (IGAN) WHO ARE AT RISK OF DISEASE PROGRESSION, BY RELEASE OF BUDESONIDE FROM THE FORMULATION
- U-3811 USE OF BIRCH TRITERPENES FOR THE TREATMENT OF WOUNDS ASSOCIATED WITH DYSTROPHIC AND JUNCTIONAL EPIDERMOLYSIS BULLOSA
- U-3812 TREATMENT OF DEPRESSION IN MDD WITH ACUTE SUICIDAL IDEATION/BEHAVIOR WITH NASALLY ADMINISTERED ESKETAMINE WITH OAD IN A PATIENT WHO HAS MISSED A DOSE IN THE MAINTENANCE PHASE AND HAD WORSENING DEPRESSION SYMPTOMS BY RETURN TO HIGHER DOSING SCHEDULE
- U-3813 TREATMENT OF TRD WITH NASALLY ADMINISTERED ESKETAMINE IN CONJUNCTION WITH AN OAD IN A PATIENT WHO HAS MISSED A DOSE DURING THE MAINTENANCE PHASE AND HAD WORSENING OF DEPRESSION SYMPTOM BY RETURNING TO HIGHER DOSING SCHEDULE
- U-3814 USE OF SUSTAINED OR EXTENDED RELEASE ORAL 25-HYDROXYVITAMIN D3 IN TREATING SECONDARY HYPERPARATHYROIDISM IN ADULT PATIENTS HAVING CHRONIC KIDNEY DISEASE STAGE 3 OR 4
- U-3815 TREATING SECONDARY HYPERPARATHYROIDISM IN STAGE 3 OR 4 CKD WITH SUSTAINED RELEASE 25-HYDROXYVITAMIN D TO REDUCE THE PATIENT'S SERUM PARATHYROID HORMONE LEVEL WHILE AVOIDING PTH OVERSUPPRESSION
- U-3816 TREATMENT OF HELICOBACTER PYLORI INFECTION USING THE ADMINISTERED DOSAGE FORMS IN ADULTS WITH DIFFERENT BODY MASS INDEX DETERMINATIONS
- U-3817 USE AS AN ANTICOAGULANT IN PATIENTS UNDERGOING PERCUTANEOUS CORONARY INTERVENTION (PCI)
- U-3818 TREATMENT OF COMPLICATED INTRA-ABDOMINAL INFECTIONS, COMPLICATED URINARY TRACT INFECTIONS, AND HOSPITAL-ACQUIRED BACTERIAL PNEUMONIA AND VENTILATOR-ASSOCIATED BACTERIAL PNEUMONIA IN ADULT AND PEDIATRIC PATIENTS (AT LEAST 31 WEEKS GESTATIONAL AGE)

PATENT AND EXCLUSIVITY TERMS

ADB 213 of 225

PATENT USE

- U-3819 A METHOD OF PREVENTING PREGNANCY BY PROVIDING AN INTRAUTERINE SYSTEM (IUS), HOLDING AN INSERTER HANDLE WITH ONE HAND, INSERTING THE IUS INTO THE UTERUS, AND MOVING A SLIDER IN THE HANDLE TO RELEASE THE IUS WITHIN THE UTERUS
- U-3820 TREATMENT OF EOSINOPHILIC ESOPHAGITIS
- U-3821 DURING LEVOKETOCONAZOLE DOSAGE TITRATION FOR THE TREATMENT OF CUSHING'S SYNDROME IN PATIENTS WHO CONCOMITANTLY USE AN OCT2 SUBSTRATE, MONITORING THE SUBJECT FOR A DOSE LIMITING EVENT AND ADJUSTING THE DOSAGE OF THE OCT2 SUBSTRATE AS NEEDED
- U-3822 AS ADJUNCTIVE TREATMENT TO LEVODOPA/CARBIDOPA IN PATIENTS WITH PARKINSON'S DISEASE EXPERIENCING "OFF" EPISODES
- U-3823 TREATMENT OF PATIENTS WITH LOCALLY ADVANCED OR METASTATIC NON-SMALL CELL LUNG CANCER WHOSE TUMORS HAVE EPIDERMAL GROWTH FACTOR RECEPTOR EXON 19 DELETIONS OR EXON 21 L858R MUTATIONS IN COMBINATION WITH PEMETREXED AND PLATINUM-BASED CHEMOTHERAPY
- U-3824 TREATMENT OF METASTATIC PANCREATIC ADENOCARCINOMA IN COMBINATION WITH OXALIPLATIN, FLUOROURACIL, AND LEUCOVORIN
- U-3825 REDUCE THE RISK OF CARDIOVASCULAR DEATH AND WORSENING HEART FAILURE IN ADULTS WITH HEART FAILURE WITH REDUCED EJECTION FRACTION, WITHOUT TYPE II DIABETES, AND HAVING AN HBA1C OF < 5.7%
- U-3826 TREATMENT OF GEOGRAPHIC ATROPHY SECONDARY TO AGE-RELATED MACULAR DEGENERATION BY INTRAVITREAL ADMINISTRATION OF 15 MG PEGCETACOPLAN EVERY OTHER MONTH
- U-3827 TREATMENT OF GEOGRAPHIC ATROPHY SECONDARY TO AGE-RELATED MACULAR DEGENERATION BY INTRAVITREAL ADMINISTRATION OF 15 MG PEGCETACOPLAN MONTHLY
- U-3828 TREATMENT OF GEOGRAPHIC ATROPHY SECONDARY TO AGE-RELATED MACULAR DEGENERATION BY INTRAVITREAL ADMINISTRATION OF 15 MG PEGCETACOPLAN MONTHLY OR EVERY OTHER MONTH
- U-3829 TREATMENT OF CORONAVIRUS DISEASE 2019 (COVID-19) IN ADULTS AND PEDIATRIC PATIENTS (WEIGHING AT LEAST 40 KG) REQUIRING HOSPITALIZATION
- U-3830 TREATMENT OF CORONAVIRUS DISEASE 2019 (COVID-19) IN NON-HOSPITALIZED ADULTS AND PEDIATRIC PATIENTS (WEIGHING AT LEAST 40 KG)
- U-3831 TREATMENT OF CORONAVIRUS DISEASE 2019 (COVID-19) IN ADULTS AND PEDIATRIC PATIENTS (BIRTH TO < 18 YEARS OF AGE WEIGHING > 1.5 KG) REQUIRING HOSPITALIZATION
- U-3832 TREATMENT OF CORONAVIRUS DISEASE 2019 (COVID-19) IN NON-HOSPITALIZED ADULTS AND PEDIATRIC PATIENTS (BIRTH TO < 18 YEARS OF AGE WEIGHING > 1.5 KG)
- U-3833 TREATMENT OF CORONAVIRUS DISEASE 2019 (COVID-19) IN ADULTS AND PEDIATRIC PATIENTS (WEIGHING AT LEAST 40 KG) REQUIRING HOSPITALIZATION AND FOR WHOM CONCOMITANT USE OF CHLOROQUINE, OR AN ANALOG OR SALT THEREOF, IS NOT RECOMMENDED
- U-3834 TREATMENT OF CORONAVIRUS DISEASE 2019 (COVID-19) IN NON-HOSPITALIZED ADULTS AND PEDIATRIC PATIENTS (WEIGHING AT LEAST 40 KG) AND FOR WHOM CONCOMITANT USE OF CHLOROQUINE, OR AN ANALOG OR SALT THEREOF, IS NOT RECOMMENDED
- U-3835 TREATMENT OF CORONAVIRUS DISEASE 2019 (COVID-19) IN ADULTS AND PEDIATRIC PATIENTS (BIRTH TO < 18 YEARS OF AGE WEIGHING > 1.5 KG) REQUIRING HOSPITALIZATION AND FOR WHOM CONCOMITANT USE OF CHLOROQUINE, OR AN ANALOG OR SALT THEREOF, IS NOT RECOMMENDED
- U-3836 TREATMENT OF CORONAVIRUS DISEASE 2019 (COVID-19) IN NON-HOSPITALIZED ADULTS AND PEDIATRIC PATIENTS (BIRTH TO < 18 YEARS OF AGE WEIGHING > 1.5 KG) AND FOR WHOM CONCOMITANT USE OF CHLOROQUINE, OR AN ANALOG OR SALT THEREOF, IS NOT RECOMMENDED
- U-3837 TREATMENT OF CORONAVIRUS DISEASE 2019 (COVID-19) IN ADULTS AND PEDIATRIC PATIENTS (WEIGHING AT LEAST 40 KG) REQUIRING HOSPITALIZATION AND FOR WHOM CONCOMITANT USE OF CHLOROQUINE, OR AN ANALOG OR SALT THEREOF, IF NOT RECOMMENDED
- U-3838 TREATMENT OF CORONAVIRUS DISEASE 2019 (COVID-19) IN NON-HOSPITALIZED ADULTS AND PEDIATRIC PATIENTS (WEIGHING AT LEAST 40 KG) AND FOR WHOM CONCOMITANT USE OF CHLOROQUINE, OR AN ANALOG OR SALT THEREOF, IF NOT RECOMMENDED
- U-3839 A METHOD OF TREATING PAIN BY ADMINISTERING BUPIVACAINE VIA LOCAL INFILTRATION IN PEDIATRIC PATIENTS AGED 6 TO LESS THAN 12 YEARS OLD UNDERGOING CARDIAC SURGERY
- U-3840 A METHOD OF TREATING PAIN BY ADMINISTERING BUPIVACAINE VIA LOCAL INFILTRATION IN PEDIATRIC PATIENTS AGED 6 TO LESS THAN 17 YEARS OLD UNDERGOING SPINE SURGERY
- U-3841 A METHOD OF ADMINISTERING BUPIVACAINE TO PRODUCE REGIONAL ANALGESIA VIA SCIATIC NERVE BLOCK IN THE POPLITEAL FOSSA IN ADULTS
- U-3842 TREATMENT OF ADULT PATIENTS WITH CHRONIC LYMPHOCYTIC LEUKEMIA (CLL) WITH 17P

PATENT AND EXCLUSIVITY TERMS

ADB 214 of 225

PATENT USE

DELETION

- U-3843 TREATMENT OF ADULT PATIENTS WITH SMALL LYMPHOCYTIC LYMPHOMA (SLL)
- U-3844 TREATMENT OF ADULT PATIENTS WITH SMALL LYMPHOCYTIC LYMPHOMA (SLL) WITH 17P DELETION
- U-3845 TREATMENT OF ADULT PATIENTS WITH WALDENSTROM'S MACROGLOBULINEMIA (WM)
- U-3846 TREATMENT OF ADULT PATIENTS WITH RELAPSED OR REFRACTORY CHRONIC LYMPHOCYTIC LEUKEMIA
- U-3847 TREATMENT OF ADULT PATIENTS WITH RELAPSED OR REFRACTORY CHRONIC LYMPHOCYTIC LEUKEMIA WITH 17P DELETION
- U-3848 TREATMENT OF ADULT PATIENTS WITH RELAPSED OR REFRACTORY SMALL LYMPHOCYTIC LYMPHOMA (SLL) WITH 17P DELETION
- U-3849 TREATMENT OF ADULT PATIENTS WITH RELAPSED OR REFRACTORY SMALL LYMPHOCYTIC LYMPHOMA (SLL)
- U-3850 TREATMENT OF CHOLESTATIC PRURITUS IN PATIENTS WITH PROGRESSIVE FAMILIAL INTRAHEPATIC CHOLESTASIS (PFIC)
- U-3851 USE OF EXBLIFEP (CEFEPIME AND ENMETAZOBACTAM) FOR TREATING COMPLICATED URINARY TRACT INFECTIONS (CUTI) INCLUDING PYELONEPHRITIS CAUSED BY DESIGNATED SUSCEPTIBLE MICROORGANISMS
- U-3852 USE OF SPECIFIED POLYMORPHS OF EXBLIFEP (CEFEPIME AND ENMETAZOBACTAM) FOR TREATING COMPLICATED URINARY TRACT INFECTIONS (CUTI) INCLUDING PYELONEPHRITIS CAUSED BY DESIGNATED SUSCEPTIBLE MICROORGANISMS
- U-3853 METHOD FOR LOWERING CHOLESTEROL LEVEL IN A HUMAN
- U-3854 LINAGLIPTIN (5 MG DAILY DOSE) AND METFORMIN (WITH OR WITHOUT INSULIN) FOR TREATING TYPE 2 DIABETES PATIENTS WITH RENAL IMPAIRMENT AND INSUFFICIENT GLYCEMIC CONTROL DESPITE PREVIOUS TREATMENT WITH METFORMIN ALONE OR IN COMBINATION WITH INSULIN
- U-3855 FOR CHRONIC WEIGHT MANAGEMENT IN ADULTS WITH AN INITIAL BODY MASS INDEX (BMI) OF: 30 KG/M² OR GREATER (OBESITY), OR 27 KG/M² OR GREATER (OVERWEIGHT) IN THE PRESENCE OF AT LEAST ONE WEIGHT-RELATED COMORBID CONDITION
- U-3856 TREATMENT OF ADULT PATIENTS WITH MANTLE CELL LYMPHOMA (MCL) WHO HAVE RECEIVED AT LEAST ONE PRIOR THERAPY
- U-3857 TREATMENT OF ADULT PATIENTS WITH WALDENSTROM'S MACROGLOBULINEMIA (WM)
- U-3858 TREATMENT OF ADULT PATIENTS WITH RELAPSED OR REFRACTORY MARGINAL ZONE LYMPHOMA (MZL) WHO HAVE RECEIVED AT LEAST ONE PRIOR ANTI-CD20-BASED REGIMEN
- U-3859 TREATMENT OF ADULT PATIENTS WITH CHRONIC LYMPHOCYTIC LEUKEMIA (CLL)
- U-3860 TREATMENT OF ADULT PATIENTS WITH RELAPSED OR REFRACTORY FOLLICULAR LYMPHOMA (FL), IN COMBINATION WITH OBINUTUZUMAB, AFTER TWO OR MORE LINES OF SYSTEMIC THERAPY
- U-3861 TREATMENT OF ADULTS WITH NONCIRRHOTIC NONALCOHOLIC STEATOHEPATITIS (NASH) WITH MODERATE TO ADVANCED LIVER FIBROSIS (CONSISTENT WITH STAGES F2 TO F3 FIBROSIS)
- U-3862 METHOD OF TREATING FRIEDRICH'S ATAXIA BY ACTIVATING THE NRF2 PATHWAY
- U-3863 METHOD OF TREATING FRIEDRICH'S ATAXIA BY ACTIVATING THE NRF2 PATHWAY WHICH REDUCES OXIDATIVE STRESS
- U-3864 METHOD FOR REDUCING A RISK OF AT LEAST ONE CARDIOVASCULAR EVENT
- U-3865 TREATMENT OF NON-24-HOUR SLEEP-WAKE DISORDER BY ADMINISTERING TASIMELTEON
- U-3866 TREATMENT OF HELICOBACTER PYLORI INFECTION IN ADULTS USING SPECIFIED DOSAGE FORM
- U-3867 METHOD FOR REDUCING A COMPOSITE ENDPOINT RISK OF MYOCARDIAL INFARCTION (MI), STROKE, CORONARY REVASCULARIZATION, UNSTABLE ANGINA REQUIRING HOSPITALIZATION, CARDIAC ARREST, AND CARDIOVASCULAR DEATH
- U-3868 METHOD FOR REDUCING ACUTE MYOCARDIAL INFARCTION RISK
- U-3869 ADJUNCTIVE TREATMENT TO LEVODOPA/CARBIDOPA IN PATIENTS WITH PARKINSON'S DISEASE EXPERIENCING OFF EPISODES
- U-3870 ADJUNCTIVE TREATMENT TO LEVODOPA/CARBIDOPA IN PATIENTS WITH PARKINSON'S DISEASE EXPERIENCING OFF EPISODES BY INCREASING L-DOPA AMOUNTS THAT REACH THE BRAIN
- U-3871 ADJUNCTIVE TREATMENT TO LEVODOPA/CARBIDOPA IN PATIENTS WITH PARKINSON'S DISEASE EXPERIENCING OFF EPISODES BY INHIBITING COMT IN THE PERIPHERY

PATENT AND EXCLUSIVITY TERMS

ADB 215 of 225

PATENT USE

- U-3872 ADJUNCTIVE TREATMENT TO LEVODOPA/CARBIDOPA IN PATIENTS WITH PARKINSON'S DISEASE EXPERIENCING OFF EPISODES BY REDUCING O-METHYLATION OF L-DOPA
- U-3873 A METHOD OF LOWERING LOW-DENSITY LIPOPROTEIN CHOLESTEROL (LDL-C) IN A HUMAN PATIENT IN NEED THEREOF COMPRISING ADMINISTRATION OF BEMPEDOIC ACID ALONE OR IN COMBINATION WITH OTHER LIPID LOWERING THERAPIES
- U-3874 IN COMBINATION WITH OTHER ANTIRETROVIRAL AGENTS FOR THE TREATMENT OF HIV-1 INFECTION IN ANTIRETROVIRAL TREATMENT-NAIVE PATIENTS 2 YEARS OF AGE AND OLDER, WEIGHING AT LEAST 14KG, WITH HIV-1 RNA LESS THAN OR EQUAL TO 100,000 AT THE START OF THERAPY
- U-3875 TREATMENT OF ADULT PATIENTS WITH RELAPSED OR REFRACTORY FOLLICULAR LYMPHOMA (FL) RECEIVING A MODERATE CYP3A INDUCER, IN COMBINATION WITH OBINUTUZUMAB, AFTER TWO OR MORE LINES OF SYSTEMIC THERAPY
- U-3876 TREATMENT OF ANEMIA DUE TO CHRONIC KIDNEY DISEASE IN ADULTS WHO HAVE BEEN RECEIVING DIALYSIS FOR AT LEAST THREE MONTHS
- U-3877 TREATMENT OF HYPERTENSION IN COMBINATION WITH OTHER ANTIHYPERTENSIVE DRUGS, INCLUDING AN ANGIOTENSIN CONVERTING ENZYME INHIBITOR, TO LOWER BLOOD PRESSURE IN ADULT PATIENTS WHO ARE NOT ADEQUATELY CONTROLLED ON OTHER DRUGS
- U-3878 TREATMENT OF HYPERTENSION IN COMBINATION WITH OTHER ANTIHYPERTENSIVE DRUGS, TO LOWER BLOOD PRESSURE IN ADULT PATIENTS WHO ARE NOT ADEQUATELY CONTROLLED ON OTHER DRUGS
- U-3879 TREATMENT OF HYPERTENSION IN COMBINATION WITH OTHER ANTIHYPERTENSIVE DRUGS, INCLUDING AN ANGIOTENSIN RECEPTOR BLOCKER, TO LOWER BLOOD PRESSURE IN ADULT PATIENTS WHO ARE NOT ADEQUATELY CONTROLLED ON OTHER DRUGS
- U-3880 TREATMENT OF CHRONIC HEPATITIS B VIRUS INFECTION IN ADULTS AND PEDIATRIC PATIENTS 6 YEARS OF AGE AND OLDER AND WEIGHING AT LEAST 25 KG
- U-3881 USE OF THE COMBINATION OF MACITENTAN AND TADALAFIL FOR THE CHRONIC TREATMENT OF ADULTS WITH PULMONARY ARTERIAL HYPERTENSION
- U-3882 METHOD OF TREATING PULMONARY ARTERIAL HYPERTENSION COMPRISING ADMINISTERING A COMBINATION OF MACITENTAN AND TADALAFIL
- U-3883 A METHOD OF LOWERING LDL-C OR REDUCING THE RISK OF CARDIOVASCULAR DISEASE IN PATIENTS WITH FAMILIAL HYPERCHOLESTEROLEMIA USING 180 MG BEMPEDOIC ACID AND 10 MG EZETIMIBE
- U-3884 A METHOD OF LOWERING LDL-C OR REDUCING THE RISK OF CARDIOVASCULAR DISEASE IN PATIENTS WITH FAMILIAL HYPERCHOLESTEROLEMIA USING A FIXED-DOSE COMBINATION OF 180 MG BEMPEDOIC ACID AND 10 MG EZETIMIBE
- U-3885 A METHOD FOR THE TREATMENT OF DUCHENNE MUSCULAR DYSTROPHY (DMD) USING GIVINOSTAT
- U-3886 METHOD OF TREATING BIPOLAR DISORDER BY ADMINISTERING A BILAYER TABLET COMPRISING OLANZAPINE AND SAMIDORPHAN
- U-3887 METHOD OF TREATING SCHIZOPHRENIA BY ADMINISTERING A BILAYER TABLET COMPRISING OLANZAPINE AND SAMIDORPHAN
- U-3888 TREATMENT OF PATIENTS WITH METASTATIC CASTRATION SENSITIVE PROSTATE CANCER
- U-3889 TREATMENT OF RELAPSING FORMS OF MULTIPLE SCLEROSIS (MS), TO INCLUDE CLINICALLY ISOLATED SYNDROME, RELAPSING-REMITTING DISEASE, AND ACTIVE SECONDARY PROGRESSIVE DISEASE, IN ADULTS RECEIVING OR WHO MAY RECEIVE A BETA-BLOCKER TREATMENT
- U-3890 A METHOD COMPRISING ADMINISTERING PEGULICIANINE TO A HUMAN AND OBTAINING AN IMAGE OF A TUMOR BED AFTER TUMOR RESECTION TO DISTINGUISH IN SITU CANCER CELLS FROM HEALTHY CELLS
- U-3891 TREATMENT OF RELAPSING FORMS OF MULTIPLE SCLEROSIS (MS), TO INCLUDE CLINICALLY ISOLATED SYNDROME, RELAPSING-REMITTING DISEASE, AND ACTIVE SECONDARY PROGRESSIVE DISEASE, IN ADULTS BEING TREATED WITH A BETA-BLOCKER OR IN ADULTS REINITIATING TREATMENT
- U-3892 TREATMENT OF EXCESSIVE DAYTIME SLEEPINESS BY ADMINISTERING SOLRIAMFETOL TO A SUBJECT HAVING A HISTORY OF BIPOLAR DISORDERS AND MODERATE OR SEVERE RENAL IMPAIRMENT
- U-3893 TREATMENT OF SICKLE CELL DISEASE BY ADMINISTERING VOXELOTOR, AS RECITED IN CLAIM 10
- U-3894 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGE 6 AND OLDER WHO ARE HOMOZYGOUS FOR THE F508DEL MUTATION OR HAVE AT LEAST ONE CFTR GENE MUTATION THAT IS RESPONSIVE TO TEZACAFTOR/IVACAFTOR WITH A COMPOSITION ACCORDING TO CLAIM 1 OF US 11951212

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-3895 TREATMENT OF PAROXYSMAL NOCTURNAL HEMOGLOBINURIA (PNH) BY ADMINISTRATION OF 200 MG OF IPTACOPAN TWICE DAILY

U-3896 TREATMENT OF HEART FAILURE WITH ORAL PELLETS

U-3897 TREATMENT OF GASTROINTESTINAL STROMAL TUMORS IN PATIENTS BEING TREATED CONCURRENTLY WITH ITRACONAZOLE

U-3898 USE OF A PHARMACEUTICAL COMPOSITION OF ABOUT 0.5M SODIUM THIOSULFATE, ABOUT 0.004M BORIC ACID, AND A PH OF BETWEEN ABOUT 6.5 AND 8.9 FOR REDUCING OTOTOXICITY IN A PEDIATRIC PATIENT RECEIVING CISPLATIN FOR THE TREATMENT OF LOCALIZED CANCER

U-3899 ADULT AND PEDIATRIC PATIENTS WEIGHING AT LEAST 10 KG AS A SOURCE OF ZINC, COPPER, MANGANESE, AND SELENIUM FOR PARENTERAL NUTRITION WHEN ORAL OR ENTERAL NUTRITION IS NOT POSSIBLE, INSUFFICIENT, OR CONTRAINDICATED

U-3900 NEONATAL AND PEDIATRIC PATIENTS WEIGHING LESS THAN 10 KG AS A SOURCE OF ZINC, COPPER, MANGANESE, AND SELENIUM FOR PARENTERAL NUTRITION WHEN ORAL OR ENTERAL NUTRITION IS NOT POSSIBLE, INSUFFICIENT, OR CONTRAINDICATED

U-3901 TREATMENT OF METASTATIC CASTRATION SENSITIVE PROSTATE CANCER

U-3902 TREATMENT OF ADULTS WITH LOCALLY ADVANCED OR METASTATIC UROTHELIAL CARCINOMA WITH SUSCEPTIBLE FGFR3 GENETIC ALTERATIONS WHOSE DISEASE HAS PROGRESSED ON OR AFTER PRIOR PD-1 OR PD-L1 INHIBITOR THERAPY

U-3903 A METHOD FOR TREATING A PEDIATRIC PATIENT WHO IS EXPERIENCING IDIOPATHIC PULMONARY ARTERIAL HYPERTENSION RESULTING IN OXYGEN DESATURATION, WHEREIN THE PATIENT IS NOT DEPENDENT ON RIGHT-TO-LEFT SHUNTING OF BLOOD

U-3904 ACUTE EXACERBATIONS OF MULTIPLE SCLEROSIS

U-3905 ALLERGIC CONJUNCTIVITIS

U-3906 ANTERIOR SEGMENT INFLAMMATION

U-3907 AS ADJUNCTIVE THERAPY FOR SHORT-TERM ADMINISTRATION (TO TIDE THE PATIENT OVER AN ACUTE EPISODE OR EXACERBATION) IN: ACUTE GOUTY ARTHRITIS

U-3908 AS ADJUNCTIVE THERAPY FOR SHORT-TERM ADMINISTRATION (TO TIDE THE PATIENT OVER AN ACUTE EPISODE OR EXACERBATION) IN: ANKYLOSING SPONDYLITIS

U-3909 AS ADJUNCTIVE THERAPY FOR SHORT-TERM ADMINISTRATION (TO TIDE THE PATIENT OVER AN ACUTE EPISODE OR EXACERBATION) IN: JUVENILE RHEUMATOID ARTHRITIS

U-3910 AS ADJUNCTIVE THERAPY FOR SHORT-TERM ADMINISTRATION (TO TIDE THE PATIENT OVER AN ACUTE EPISODE OR EXACERBATION) IN: PSORIATIC ARTHRITIS

U-3911 AS ADJUNCTIVE THERAPY FOR SHORT-TERM ADMINISTRATION (TO TIDE THE PATIENT OVER AN ACUTE EPISODE OR EXACERBATION) IN: RHEUMATOID ARTHRITIS

U-3912 ATOPIC DERMATITIS

U-3913 CHORIORETINITIS

U-3914 DIFFUSE POSTERIOR UVEITIS AND CHOROIDITIS

U-3915 DURING AN EXACERBATION OR AS MAINTENANCE THERAPY IN SELECTED CASES OF: SYSTEMIC DERMATOMYOSITIS (POLYMYOSITIS)

U-3916 DURING AN EXACERBATION OR AS MAINTENANCE THERAPY IN SELECTED CASES OF: SYSTEMIC LUPUS ERYTHEMATOSUS

U-3917 IRIDOCYCLITIS

U-3918 IRITIS

U-3919 KERATITIS

U-3920 OPTIC NEURITIS

U-3921 SERUM SICKNESS

U-3922 SEVERE ACUTE AND CHRONIC ALLERGIC AND INFLAMMATORY PROCESSES INVOLVING THE EYE AND ITS ADNEXA

U-3923 SEVERE ERYTHEMA MULTIFORME (STEVENS-JOHNSON SYNDROME)

U-3924 SEVERE PSORIASIS

U-3925 SYMPTOMATIC SARCOIDOSIS

U-3926 TO INDUCE A DIURESIS OR A REMISSION OF PROTEINURIA IN THE NEPHROTIC SYNDROME WITHOUT UREMIA OF THE IDIOPATHIC TYPE OR THAT DUE TO LUPUS ERYTHEMATOSUS

PATENT AND EXCLUSIVITY TERMS

ADB 217 of 225

PATENT USE

- U-3927 USE TO INCREASE LINEAR GROWTH IN PEDIATRIC PATIENTS WITH ACHONDROPLASIA WITH OPEN EPIPHYSES
- U-3928 FOR THE TREATMENT OF INTERMEDIATE OR HIGH-RISK MYELOFIBROSIS IN SUBJECTS HAVING A BASELINE PLATELET COUNT OF LESS THAN 50 BILLION/L
- U-3929 MAINTENANCE TREATMENT WITH 15 MG/KG OF BODY WEIGHT BEVACIZUMAB EVERY THREE WEEKS OF ADVANCED EPITHELIAL OVARIAN CANCER IN COMPLETE OR PARTIAL RESPONSE TO FIRST-LINE PLATINUM-BASED CHEMOTHERAPY AND ASSOCIATED WITH HRD-POSITIVE STATUS
- U-3930 MAINTENANCE TREATMENT WITH 15 MG/KG OF BODY WEIGHT BEVACIZUMAB EVERY THREE WEEKS OF FALLOPIAN TUBE CANCER IN COMPLETE OR PARTIAL RESPONSE TO FIRST-LINE PLATINUM-BASED CHEMOTHERAPY AND ASSOCIATED WITH HRD-POSITIVE STATUS
- U-3931 MAINTENANCE TREATMENT WITH 15 MG/KG OF BODY WEIGHT BEVACIZUMAB EVERY THREE WEEKS OF PRIMARY PERITONEAL CANCER IN COMPLETE OR PARTIAL RESPONSE TO FIRST-LINE PLATINUM-BASED CHEMOTHERAPY AND ASSOCIATED WITH HRD-POSITIVE STATUS
- U-3932 TREATMENT OF PATIENTS 12 YEARS AND OLDER WITH WARTS, HYPOGAMMAGLOBULINEMIA, INFECTIONS, AND MYELOKATHEXIS (WHIM) SYNDROME
- U-3933 TREATMENT OF EXTRAVASCULAR HEMOLYSIS (EVH) IN ADULTS WITH PAROXYSMAL NOCTURNAL HEMOGLOBINURIA (PNH) AS ADD-ON THERAPY TO RAVULIZUMAB OR ECULIZUMAB
- U-3934 TREATMENT OF NARCOLEPSY-RELATED CATAPLEXY OR EDS IN A HUMAN PATIENT WITH A ONCE-NIGHTLY GHB FORMULATION BY INITIATING TREATMENT WITH A DOSE EQUIVALENT TO 4.5 G OF SODIUM OXYBATE AND UPTITRATING IN INCREMENTSEQUIVALENT TO 1.5 G OF SODIUM OXYBATE
- U-3935 ACUTE TREATMENT OF AGITATION ASSOCIATED WITH SCHIZOPHRENIA OR BIPOLAR I OR II DISORDER BY SUBLINGUAL OR BUCCAL ADMINISTRATION
- U-3936 TREATMENT OF ADULT PATIENTS WITH SEVERE APLASTIC ANEMIA WHO HAVE HAD AN INSUFFICIENT RESPONSE TO IMMUNOSUPPRESSIVE THERAPY
- U-3937 TREATMENT OF THROMBOCYTOPENIA IN ADULT PATIENTS WITH CHRONIC HEPATITIS C TO ALLOW THE INITIATION AND MAINTENANCE OF INTERFERON-BASED THERAPY
- U-3938 TREATMENT OF THROMBOCYTOPENIA IN ADULT AND PEDIATRIC PATIENTS 6 YEARS AND OLDER WITH PERSISTENT OR CHRONIC IMMUNE THROMBOCYTOPENIA (ITP)
- U-3939 INTRAMUSCULAR ADMINISTRATION OF RISPERIDONE IN EXTENDED RELEASE INJECTABLE SUSPENSION FOR TREATMENT OF SCHIZOPHRENIA IN ADULTS
- U-3940 TREATMENT OF BIPOLAR DISORDER OR BIPOLAR DEPRESSION, INCLUDING ASSOCIATED DEPRESSIVE EPISODES
- U-3941 ADMINISTRATION OF RISPERIDONE IN EXTENDED RELEASE INJECTABLE SUSPENSION FOR TREATMENT OF SCHIZOPHRENIA IN ADULTS
- U-3942 TREATMENT OF SCHIZOPHRENIA IN ADULTS BY INTRAMUSCULAR ADMINISTRATION OF EXTENDED RELEASE INJECTABLE COMPOSITION
- U-3943 TREATMENT OF SCHIZOPHRENIA IN ADULTS BY ADMINISTRATION OF EXTENDED RELEASE INJECTABLE COMPOSITION
- U-3944 METHOD FOR DELIVERING AN EFFECTIVE AMOUNT OF CLONIDINE FOR A 24-HOUR PERIOD USING A SINGLE ORAL CLONIDINE COMPOSITION ACCORDING TO CLAIM 1 PRIOR TO BED TIME
- U-3945 TREATMENT OF PATIENTS 2 YEARS OF AGE AND OLDER WITH ACTIVE POLYARTICULAR JUVENILE IDIOPATHIC ARTHRITIS WHO HAVE HAD AN INADEQUATE RESPONSE OR INTOLERANCE TO ONE OR MORE TNF BLOCKERS
- U-3946 TREATMENT OF PEDIATRIC PATIENTS 2 YEARS OF AGE AND OLDER WITH ACTIVE PSORIATIC ARTHRITIS WHO HAVE HAD AN INADEQUATE RESPONSE OR INTOLERANCE TO ONE OR MORE TNF BLOCKERS
- U-3947 TOPICAL TREATMENT OF ROSACEA
- U-3948 USE OF A PHARMACEUTICAL COMPOSITION HAVING ABOUT 0.5M AQUEOUS ANHYDROUS SODIUM THIOSULFATE AND A CONCENTRATION OF BORATE IONS OF LESS THAN 0.05% FOR REDUCING OTOTOXICITY IN A PEDIATRIC PATIENT RECEIVING CISPLATIN FOR TREATMENT OF LOCALIZED CANCER
- U-3949 ADULT AND PEDIATRIC PATIENTS 2 YEARS OF AGE AND OLDER WITH LOCALLY ADVANCED OR METASTATIC SOLID TUMORS WITH A RET GENE FUSION THAT HAS PROGRESSED ON OR FOLLOWING PRIOR SYSTEMIC TREATMENT OR WHO HAVE NO SATISFACTORY ALTERNATIVE TREATMENT OPTIONS
- U-3950 TREATMENT OF ADULT OR PEDIATRIC PATIENTS 2 YEARS OF AGE OR OLDER WITH ADVANCED OR METASTATIC MEDULLARY THYROID CANCER (MTC) WITH A RET MUTATION, AS DETECTED BY AN FDA-APPROVED TEST, WHO REQUIRE SYSTEMIC THERAPY

PATENT AND EXCLUSIVITY TERMS

ADB 218 of 225

PATENT USE

- U-3951 TREATMENT OF ADULT OR PEDIATRIC PATIENTS 2 YEARS OF AGE OR OLDER WITH ADVANCED OR METASTATIC THYROID CANCER WITH A RET GENE FUSION, AS DETECTED BY AN FDA-APPROVED TEST, WHO REQUIRE SYSTEMIC THERAPY AND WHO ARE RADIOACTIVE IODINE-REFRACTORY
- U-3952 USE OF A PHARMACEUTICAL COMPOSITION HAVING A CONCENTRATION OF ABOUT 0.5M AQUEOUS ANHYDROUS SODIUM THIOSULFATE AND ABOUT 0.004M BORIC ACID FOR REDUCING OTOTOXICITY IN A PEDIATRIC PATIENT RECEIVING CISPLATIN FOR TREATMENT OF LOCALIZED CANCER
- U-3953 TREATMENT, IN COMBINATION WITH CETUXIMAB, OF ADULT PATIENTS WITH KRAS G12C-MUTED LOCALLY ADVANCED OR METASTATIC COLORECTAL CANCER, PER FDA APPROVED TEST, WHERE PRIOR TREATMENT WITH FLUOROPYRIMIDINE-, OXALIPLATIN-, AND IRINOTECAN-BASED CHEMOTHERAPY
- U-3954 ADJUNCTIVE THERAPY IN THE TREATMENT OF PRIMARY GENERALIZED TONIC-CLONIC SEIZURES IN ADULTS AND IN PEDIATRIC PATIENTS WEIGHING AT LEAST 50 KG
- U-3955 TREATMENT OF PRIMARY BILIARY CHOLANGITIS (PBC) IN COMBINATION WITH URSODEOXYCHOLIC ACID (UDCA) IN ADULTS WHO HAVE HAD AN INADEQUATE RESPONSE TO UDCA, OR AS MONOTHERAPY IN PATIENTS UNABLE TO TOLERATE UDCA
- U-3956 TREATMENT OF PATIENTS WITH MYELODYSPLASTIC SYNDROMES (MDS) WITH TRANSFUSION-DEPENDENT ANEMIA
- U-3957 TREATMENT OF GASTROINTESTINAL STROMAL TUMOR COMPRISING ADMINISTERING ONCE DAILY, ONE OR MORE TABLETS COMPRISING RIPRETINIB
- U-3958 TREATMENT OF ADVANCED GASTROINTESTINAL STROMAL TUMOR IN PATIENTS PREVIOUSLY ADMINISTERED 4 LINES OF PRIOR THERAPY, WHERE THE FIRST LINE OF PRIOR THERAPY IS IMATINIB
- U-3959 TREATMENT OF GASTROINTESTINAL STROMAL TUMORS IN A PATIENT SUFFERING FROM GRADE 2 ARTHRALGIA WHILE BEING ADMINISTERED RIPRETINIB DAILY
- U-3960 TREATMENT OF ADVANCED GASTROINTESTINAL STROMAL TUMOR IN PATIENTS PREVIOUSLY ADMINISTERED IMATINIB
- U-3961 TREATMENT OF ADULT AND PEDIATRIC PATIENTS >12 YEARS WITH SOLID TUMORS AND NTRK GENE FUSION THAT ARE LOCALLY ADVANCED OR METASTATIC OR LIKELY SURGICALLY UNRESECTABLE, AND HAVE PROGRESSED FOLLOWING TREATMENT OR HAVE NO SATISFACTORY ALTERNATIVE THERAPY
- U-3962 FOR THE MAINTENANCE TREATMENT OF CHRONIC OBSTRUCTIVE PULMONARY DISEASE (COPD) IN ADULT PATIENTS
- U-3963 TREATMENT OF CLINICALLY IMPORTANT HYPOTENSION OCCURRING IN THE SETTING OF ANESTHESIA
- U-3964 METHODS OF ADMINISTERING TESTOSTERONE UNDECANOATE FOR RESTORING TESTOSTERONE CONCENTRATION IN A HYPOGONADAL MALE SUBJECT
- U-3965 METHODS OF CONTINUING OR DISCONTINUING A DOSAGE REGIMEN OF A TESTOSTERONE UNDECANOATE PHARMACEUTICAL COMPOSITION TO A HYPOGONADAL MALE SUBJECT
- U-3966 REDUCTION OF HETEROTOPIC OSSIFICATION IN PATIENTS WITH FIBRODYSPLASIA OSSIFICANS (MYOSITIS) PROGRESSIVA
- U-3967 TREATING AN ACUTE BACTERIAL SKIN AND SKIN STRUCTURE INFECTION (ABSSSI) IN AN OVERWEIGHT OR OBESE PATIENT BY INTRAVENOUSLY (IV) ADMINISTERING 300MG OF DELAFLOXACIN OR A PHARMACEUTICALLY ACCEPTABLE SALT, TWICE A DAY
- U-3968 DOSING REGIMEN FOR THE TREATMENT OF SCHIZOPHRENIA IN ADULTS BY ADMINISTERING AN INITIAL DOSE OF PALIPERIDONE PALMITATE FOLLOWED BY MONTHLY DOSE(S)
- U-3969 DOSING REGIMEN FOR THE TREATMENT OF SCHIZOAFFECTIVE DISORDER IN ADULTS AS MONOTHERAPY AND AS AN ADJUNCT TO MOOD STABILIZERS OR ANTIDEPRESSANTS BY ADMINISTERING AN INITIAL DOSE OF PALIPERIDONE PALMITATE FOLLOWED BY MONTHLY DOSE(S)
- U-3970 TOPICAL TREATMENT OF MILD TO MODERATE ATOPIC DERMATITIS
- U-3971 TREATMENT OF GASTROINTESTINAL STROMAL TUMORS IN PATIENTS SUFFERING FROM PALMAR-PLANTAR ERYTHRODYSESTHESIA SYNDROME
- U-3972 TREATMENT OF ADULTS WITH LOCALLY ADVANCED OR METASTATIC UROTHELIAL CARCINOMA WITH SUSCEPTIBLE FGFR3 GENETIC ALTERATIONS (Y373C IDENTIFIED) WHOSE DISEASE HAS PROGRESSED ON OR AFTER AT LEAST ONE LINE OF PRIOR SYSTEMIC THERAPY
- U-3973 TREATMENT OF CHOLESTATIC PRURITUS IN PATIENTS 12 MONTHS OF AGE AND OLDER WITH PROGRESSIVE FAMILIAL INTRAHEPATIC CHOLESTASIS (PFIC)

PATENT AND EXCLUSIVITY TERMS

ADB 219 of 225

PATENT USE

- U-3974 TREATMENT OF CHOLESTATIC PRURITUS IN PATIENTS 3 MONTHS OF AGE AND OLDER WITH ALAGILLE SYNDROME (ALGS)
- U-3975 IN COMBINATION WITH FULVESTRANT AS INITIAL ENDOCRINE-BASED THERAPY OR FOLLOWING DISEASE PROGRESSION ON ENDOCRINE THERAPY, FOR THE TREATMENT OF ADULT PATIENTS WITH HR-POSITIVE, HER2-NEGATIVE ADVANCED OR METASTATIC BREAST CANCER
- U-3976 TREATMENT OF ADULT PATIENTS WITH ALOPECIA AREATA
- U-3977 A METHOD OF TREATING A GLIOMA CHARACTERIZED BY AN IDH1 MUTATION FOLLOWING SURGERY, WHEREIN THE GLIOMA IS GRADE 2 ASTROCYTOMA OR OLIGODENDROGLIOMA
- U-3978 A METHOD OF TREATING A CANCER CHARACTERIZED BY AN IDH1 OR IDH2 MUTATION OR COMBINATION THEREOF FOLLOWING SURGERY, WHEREIN THE CANCER IS GRADE 2 ASTROCYTOMA OR OLIGODENDROGLIOMA
- U-3979 NASAL ADMINISTRATION OF EPINEPHRINE FOR THE TREATMENT OF ALLERGIC REACTIONS (TYPE I), INCLUDING ANAPHYLAXIS
- U-3980 TREATMENT OF PRIMARY IMMUNOGLOBULIN A NEPHROPATHY (IGAN)
- U-3981 TREATMENT OF THE FOLLOWING INFECTIONS: COMPLICATED SKIN AND SKIN STRUCTURE INFECTIONS AND STAPHYLOCOCCUS AUREUS BLOOD STREAM INFECTIONS
- U-3982 TREATMENT OF HYPOPARATHYROIDISM IN ADULTS
- U-3983 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 1 YEAR TO 5 YEARS WHO ARE HOMOZYGOUS FOR THE F508DEL CFTR GENE MUTATION COMPRISING ADMINISTERING AN EFFECTIVE AMOUNT OF IVA AND FORM I LUM AS RECITED IN, E.G., CLAIM 1 OF US 12065432
- U-3984 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 6 YEARS AND OLDER WHO ARE HOMOZYGOUS FOR THE F508DEL CFTR GENE MUTATION COMPRISING ADMINISTERING AN EFFECTIVE AMOUNT OF IVA AND FORM I LUM AS RECITED IN, E.G., CLAIM 1 OF US 12065432
- U-3985 FIRST-LINE TREATMENT OF ADULTS WITH LOCALLY ADVANCED OR METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) WITH EGFR EXON 19 DELETIONS OR EXON 21 L858R SUBSTITUTION MUTATIONS, IN COMBINATION WITH AMIVANTAMAB
- U-3986 A METHOD OF TREATING AGITATION ASSOCIATED WITH SCHIZOPHRENIA OR BIPOLAR I OR II DISORDER
- U-3987 TOPICAL TREATMENT OF ACNE
- U-3988 USE FOR THE TREATMENT OF FOCAL SEIZURES ASSOCIATED WITH LENNOX-GASTAUT SYNDROME
- U-3989 USE FOR THE TREATMENT OF FOCAL SEIZURES ASSOCIATED WITH TUBEROUS SCLEROSIS COMPLEX
- U-3990 TREATMENT OF HYPERTENSION IN PATIENTS WHO ARE IN NEED OF A LIQUID COMPOSITION OF TERAZOSIN
- U-3991 TREATMENT OF SYMPTOMATIC BENIGN PROSTATIC HYPERPLASIA (BPH) IN PATIENTS WHO ARE IN NEED OF A LIQUID COMPOSITION OF TERAZOSIN
- U-3992 ACUTE TREATMENT OF MIGRAINE WITH OR WITHOUT AURA IN A PATIENT UNDERGOING TREATMENT WITH A WEAK OR MODERATE CYP3A4 INDUCER
- U-3993 TREATMENT OF PRIMARY IMMUNOGLOBULIN A NEPHROPATHY (IGAN) IN ADULTS AT RISK FOR DISEASE PROGRESSION
- U-3994 METHOD OF TREATING SEVERE ALOPECIA AREATA IN ADULTS AND ADOLESCENTS 12 YEARS AND OLDER BY ADMINISTERING RITLECTINIB
- U-3995 TO RESTORE BLOOD PRESSURE IN ADULT PATIENTS WITH ACUTE HYPOTENSIVE STATES
- U-3996 FIRST-LINE TREATMENT OF ADULTS PATIENTS WITH ADVANCED RENAL CELL CARCINOMA BY ADMINISTERING LENVIMA IN COMBINATION WITH PEMBROLIZUMAB
- U-3997 TREATMENT OF PATIENTS WITH PMMR/NOT MSI-H ADVANCED ENDOMETRIAL CARCINOMA, HAVE DISEASE PROGRESSION FOLLOWING PRIOR SYSTEMIC THERAPY, AND ARE NOT CANDIDATES FOR CURATIVE SURGERY OR RADIATION, BY ADMINISTERING LENVIMA IN COMBINATION WITH PEMBROLIZUMAB
- U-3998 ADJUVANT TREATMENT OF ADULTS WITH HORMONE RECEPTOR (HR)-POSITIVE, HUMAN EPIDERMAL GROWTH FACTOR RECEPTOR 2 (HER2)-NEGATIVE STAGE II AND III EARLY BREAST CANCER AT HIGH RISK OF RECURRENCE
- U-3999 IN COMBINATION WITH AN AROMATASE INHIBITOR FOR THE ADJUVANT TREATMENT OF ADULTS WITH HORMONE RECEPTOR (HR)-POSITIVE, HUMAN EPIDERMAL GROWTH FACTOR RECEPTOR 2 (HER2)-NEGATIVE STAGE II AND III EARLY BREAST CANCER AT HIGH RISK OF RECURRENCE

PATENT AND EXCLUSIVITY TERMS

ADB 220 of 225

PATENT USE

- U-4000 TREATMENT OF DEPRESSIVE EPISODES OR MAJOR DEPRESSIVE EPISODES ASSOCIATED WITH BIPOLAR II DISORDER
- U-4001 TREATMENT OF ADVANCED PROSTATIC CANCER
- U-4002 POSITRON EMISSION TOMOGRAPHY (PET) OF PROSTATE-SPECIFIC MEMBRANE ANTIGEN (PSMA)- POSITIVE LESIONS IN MEN WITH PROSTATE CANCER
- U-4003 TREATMENT OF NOCTURIA DUE TO NOCTURNAL POLYURIA IN ADULTS WHO AWAKEN AT LEAST 2 TIMES PER NIGHT TO VOID BY INTRANASALLY ADMINISTERING A PLUME OF DROPLETS COMPRISING A DOSE OF 0.75 MCG OR 1.5 MCG DESMOPRESSIN
- U-4004 TREATMENT OF PARKINSONISM THAT MAY FOLLOW CARBON MONOXIDE INTOXICATION
- U-4005 TREATMENT OF PARKINSONISM THAT MAY FOLLOW MANGANESE INTOXICATION
- U-4006 TREATMENT OF A DISORDER TREATABLE WITH GHB USING A SINGLE DOSE OF A GHB FORMULATION IN AN AMOUNT EQUIVALENT TO 3 G TO 12 G OF SODIUM OXYBATE WHERE THE FORMULATION CONTAINS AN IMMEDIATE RELEASE PORTION AND A MODIFIED RELEASE PORTION
- U-4007 METHOD OF TREATING ACROMEGALY PATIENTS WHO HAVE HAD INADEQUATE RESPONSE TO OR CANNOT BE TREATED WITH SURGICAL RESECTION, PITUITARY IRRADIATION, AND BROMOCRIPTINE MESYLATE AT MAXIMALLY TOLERATED DOSES
- U-4008 METHOD OF TREATING PROFUSE WATERY DIARRHEA ASSOCIATED WITH VASOACTIVE INTESTINAL PEPTIDE TUMORS (VIPOMAS)-SECRETING TUMORS
- U-4009 METHOD OF TREATING SEVERE DIARRHEA AND FLUSHING EPISODES ASSOCIATED WITH METASTATIC CARCINOID TUMORS
- U-4010 TREATMENT OF PATIENTS WITH LOCALLY ADVANCED, UNRESECTABLE (STAGE III) NSCLC WHOSE DISEASE HAS NOT PROGRESSED DURING OR FOLLOWING PLATINUM-BASED CHEMORADIATION THERAPY AND WHOSE TUMORS HAVE EGFR EXON 19 DELETIONS OR EXON 21 L858R MUTATIONS
- U-4011 METHOD OF POSITRON EMISSION TOMOGRAPHY (PET) FOR CARDIAC IMAGING
- U-4012 PRETREATMENT AGAINST THE LETHAL EFFECTS OF SOMAN NERVE AGENT POISONING
- U-4013 A METHOD OF ADJUNCTIVE TREATING, PREVENTING AND/OR AMELIORATING SEIZURES IN A PERSON WITH DRAVET SYNDROME COMPRISING ADMINISTERING FENFLURAMINE OR A SALT THEREOF IN A DOSE OF 0.2 MG/KG/DAY TO 0.5/MG/KG/DAY AND DOSES OF STIRIPENTOL AND CLOBAZAM
- U-4014 TREATMENT OF CHRONIC GRAFT-VERSUS-HOST DISEASE (CHRONIC GVHD) AFTER FAILURE OF AT LEAST TWO PRIOR LINES OF SYSTEMIC THERAPY IN ADULT AND PEDIATRIC PATIENTS 12 YEARS AND OLDER WITH A GRADE 3 INFECTION ADVERSE REACTION
- U-4015 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETIS BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 1 AND 10
- U-4016 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETIS BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 1, 10, AND 11
- U-4017 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETIS BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 12 AND 21
- U-4018 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETIS BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 12, 21, AND 22
- U-4019 TREATMENT OF BIPOLAR DEPRESSION BY INHIBITING THE 5-HT2A RECEPTOR, INHIBITING THE SEROTONIN TRANSPORTER, AND/OR MODULATING DOPAMINE D2 RECEPTOR FUNCTION, OR TREATMENT OF BIPOLAR DISORDER
- U-4020 TREATMENT OF SCHIZOPHRENIA BY INHIBITING THE 5-HT2A RECEPTOR, INHIBITING THE SEROTONIN TRANSPORTER, AND/OR MODULATING DOPAMINE D2 RECEPTOR FUNCTION, OR TREATMENT OF SCHIZOPHRENIA
- U-4021 USE OF ARIMOCLOMOL, IN COMBINATION WITH MIGLUSTAT, FOR TREATMENT OF NEUROLOGICAL MANIFESTATIONS OF NIEMANN-PICK DISEASE TYPE C (NPC)
- U-4022 TREATMENT OF A DISORDER IN A HUMAN PATIENT WITH A ONCE-NIGHTLY GHB FORMULATION BY INITIATING TREATMENT WITH A DOSE EQUIVALENT TO 4.5 G OF SODIUM OXYBATE AND UPTITRATING IN INCREMENTS EQUIVALENT TO 1.5 G OF SODIUM OXYBATE
- U-4023 TREATMENT OF TYPE 2 DIABETES MELLITUS BY ADMINISTRATION OF EMPAGLIFLOZIN, LINAGLIPTIN AND METFORMIN

PATENT AND EXCLUSIVITY TERMS

ADB 221 of 225

PATENT USE

- U-4024 COMBINATION WITH PALBOCICLIB AND FULVESTRANT FOR TREATMENT OF ADULTS WITH ENDOCRINE-RESISTANT PIK3CA-MUTATED HR-POSITIVE HER2-NEGATIVE LOCALLY ADVANCED OR METASTATIC BREAST CANCER FOLLOWING RECURRENCE ON OR AFTER COMPLETING ADJUVANT ENDOCRINE THERAPY
- U-4025 A METHOD FOR THE TREATMENT OF POST-OPERATIVE INFLAMMATION FOLLOWING OCULAR SURGERY
- U-4026 USE IN TREATING UNCOMPLICATED URINARY TRACT INFECTIONS CAUSED BY ESCHERICHIA COLI, KLEBSIELLA PNEUMONIAE, OR PROTEUS MIRABILIS
- U-4027 TREATMENT OF GASTRIC ADENOCARCINOMA
- U-4028 TREATMENT OF HEAD AND NECK CANCER
- U-4029 ACUTE TREATMENT OF AGITATION IN PATIENTS WITH SEVERE HEPATIC IMPAIRMENT
- U-4030 TREATMENT OF MOTOR FLUCTUATIONS IN ADULTS WITH ADVANCED PARKINSON'S DISEASE
- U-4031 REDUCTION OF HETEROTOPIC OSSIFICATION IN PATIENTS WITH FIBRODYSPLASIS (MYOSITIS) OSSIFICANS PROGRESSIVA
- U-4032 TREATMENT OF ATTENTION DEFICIT HYPERACTIVITY DISORDER (ADHD)
- U-4033 METHOD OF PROVIDING LOCAL OR REGIONAL ANALGESIA, FOR EXAMPLE, VIA INFILTRATION TO PROVIDE POSTSURGICAL LOCAL ANALGESIA OR VIA INTERSCALENE BRACHIAL PLEXUS NERVE BLOCK TO PROVIDE POSTSURGICAL REGIONAL ANALGESIA
- U-4034 TREATMENT OF SEIZURES ASSOCIATED WITH CYCLIN-DEPENDENT KINASE-LIKE-5 (CDLK5) DEFICIENCY DISORDER (CDD) IN PATIENTS 2 YEARS OF AGE AND OLDER
- U-4035 TREATMENT OF ADULT PATIENTS WITH ADVANCED PROSTATE CANCER, MODIFIED BY DOSE-SEPARATING ADMINISTRATIONS OF RELUGOLIX AND A P-GP INHIBITOR, WHEN CO-ADMINISTRATION OF RELUGOLIX AND THE P-GP INHIBITOR IS UNAVOIDABLE
- U-4036 TREATMENT OF ADULT AND PEDIATRIC PATIENTS 12 YEARS AND OLDER WITH SOMATOSTATIN RECEPTOR-POSITIVE GASTROENTEROPANCREATIC NEUROENDOCRINE TUMORS (GEP-NETS), INCLUDING FOREGUT, MIDGUT, AND HINDGUT NEUROENDOCRINE TUMORS
- U-4037 METHOD OF TREATING PAIN, FOR EXAMPLE, VIA INFILTRATION TO PROVIDE LOCAL ANALGESIA OR VIA INTERSCALENE BRACHIAL PLEXUS NERVE BLOCK TO PROVIDE REGIONAL ANALGESIA
- U-4038 ADJUNCTIVE TREATMENT TO LEVODOPA/CARBIDOPA IN PATIENTS WITH PARKINSON'S DISEASE EXPERIENCING OFF EPISODES BY ADMINISTERING WITHOUT FOOD
- U-4039 ADULT AND PEDIATRIC PATIENTS AS A SOURCE OF SELENIUM FOR PARENTERAL NUTRITION WHEN ORAL OR ENTERAL NUTRITION IS NOT POSSIBLE, INSUFFICIENT, OR CONTRAINDICATED
- U-4040 TREATMENT OF ADULT PATIENTS WITH LOCALLY ADVANCED UNRESECTABLE OR METASTATIC MALIGNANT PERIVASCULAR EPITHELIOD CELL TUMOR (PECOMA) WITH A DOSE ADMINISTERED ON DAYS 1 AND 8 OF A 21-DAY CYCLE
- U-4041 TREATMENT OF ADULT PATIENTS WITH LOCALLY ADVANCED UNRESECTABLE OR METASTATIC MALIGNANT PERIVASCULAR EPITHELIOD CELL TUMOR (PECOMA) WITH A DOSE OF 45 MG/M2 OR 56 MG/M2 ADMINISTERED ON DAYS 1 AND 8 OF A 21-DAY CYCLE
- U-4042 TREATMENT OF ADULT PATIENTS WITH LOCALLY ADVANCED UNRESECTABLE OR METASTATIC MALIGNANT PERIVASCULAR EPITHELIOD CELL TUMOR (PECOMA) WITH A DOSE OF 45 MG/M2 ADMINISTERED ON DAYS 1 AND 8 OF A 21-DAY CYCLE
- U-4043 EMROSI IS INDICATED TO TREAT INFLAMMATORY LESIONS (PAPULES AND PUSTULES) OF ROSACEA IN ADULTS
- U-4044 TREATMENT OF IDIOPATHIC HYPERSOMNIA WITH A MIXTURE OF SODIUM, POTASSIUM, MAGNESIUM, AND CALCIUM SALTS OF GHB
- U-4045 TREATMENT OF RELAPSED OR REFRACTORY ACUTE LEUKEMIA WITH A LYSINE METHYLTRANSFERASE 2A GENE (KMT2A) TRANSLOCATION IN ADULT AND PEDIATRIC PATIENTS 1 YEAR AND OLDER
- U-4046 TREATMENT OF THE CARDIOMYOPATHY OF WILD-TYPE OR VARIANT TRANSTHYRETIN-MEDIATED AMYLOIDOSIS (ATTR-CM)
- U-4047 A METHOD FOR TREATING ULCERATIVE COLITIS BY ADMINISTERING A THERAPEUTICALLY EFFECTIVE AMOUNT OF ESTRASIMOD ARGININE AS CLAIMED
- U-4048 TOPICAL TREATMENT OF ATOPIC DERMATITIS IN ADULTS AND PEDIATRIC PATIENTS 2 YEARS OF AGE AND OLDER
- U-4049 ADJUNCTIVE TREATMENT OF CLASSIC CONGENITAL ADRENAL HYPERPLASIA (CAH)
- U-4050 USE IN REDUCING TRIGLYCERIDES IN ADULTS WITH FAMILIAL CHYLOMICRONEMIA SYNDROME (FCS)

PATENT AND EXCLUSIVITY TERMS

ADB 222 of 225

PATENT USE

- U-4051 BRAFTOVI IS A KINASE INHIBITOR INDICATED IN COMBINATION WITH CETUXIMAB AND MFOLFOX6, FOR THE TREATMENT OF PATIENTS WITH METASTATIC COLORECTAL CANCER (MCRC) WITH A BRAF V600E MUTATION, AS DETECTED BY AN FDA-APPROVED TEST
- U-4052 TREATMENT OF CF IN PATIENTS AGED 2 TO <6 YEARS WHO HAVE IN THE CFTR GENE AT LEAST ONE F508DEL MUTATION OR A RESPONSIVE MUTATION BASED ON CLINICAL AND/OR IN VITRO DATA COMPRISING ADMINISTERING A COMPOSITION ACCORDING TO CLAIM 1 OF US 11752106
- U-4053 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 2 TO <6 YEARS WHO HAVE IN THE CFTR GENE AT LEAST ONE F508DEL MUTATION OR A RESPONSIVE MUTATION BASED ON CLINICAL AND/OR IN VITRO DATA WITH AN EFFECTIVE AMOUNT OF ELX, TEZ, AND IVA
- U-4054 TREATMENT OF CF IN PATIENTS 6 YEARS AND OLDER WHO HAVE IN THE CFTR GENE AT LEAST ONE F508DEL MUTATION OR A RESPONSIVE MUTATION BASED ON CLINICAL AND/OR IN VITRO DATA BY ADMINISTERING ELX, IVA, AND A SOLID DISPERSION ACCORDING TO US 11578062 CLAIM 1
- U-4055 TREATMENT OF CF IN PATIENTS AGED 2 TO <6 YEARS WHO HAVE IN THE CFTR GENE AT LEAST ONE F508DEL MUTATION OR A RESPONSIVE MUTATION BASED ON CLINICAL AND/OR IN VITRO DATA COMPRISING ADMINISTERING THE COMPOSITION RECITED IN US 11564916 CLAIM 1
- U-4056 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 6 YEARS AND OLDER WHO HAVE IN THE CFTR GENE AT LEAST ONE F508DEL MUTATION OR A RESPONSIVE MUTATION BASED ON CLINICAL AND/OR IN VITRO DATA WITH AN EFFECTIVE AMOUNT OF ELX, TEZ, AND IVA
- U-4057 TREATMENT OF CF IN PATIENTS AGED 2 TO <6 YEARS WHO HAVE IN THE CFTR GENE AT LEAST ONE F508DEL MUTATION OR A RESPONSIVE MUTATION BASED ON CLINICAL AND/OR IN VITRO DATA BY ADMINISTERING DAILY ELX (100 MG OR 80 MG); TEZ; AND IVA
- U-4058 TREATMENT OF CF IN PATIENTS AGED 2 TO <6 YEARS WHO HAVE IN THE CFTR GENE AT LEAST ONE F508DEL MUTATION OR A RESPONSIVE MUTATION BASED ON CLINICAL AND/OR IN VITRO DATA WITH A COMPOUND OF CLAIM 1 OR COMPOSITION OF CLAIM 29 OF US 11426407
- U-4059 TREATMENT OF CF IN PATIENTS AGED 2 TO <6 YEARS WHO HAVE IN THE CFTR GENE AT LEAST ONE F508DEL MUTATION OR A RESPONSIVE MUTATION BASED ON CLINICAL AND/OR IN VITRO DATA WITH A COMPOSITION ACCORDING TO CLAIM 1 OF US 11179367
- U-4060 TREATMENT OF CF IN PATIENTS AGED 2 TO <6 YEARS WHO HAVE IN THE CFTR GENE AT LEAST ONE F508DEL MUTATION OR A RESPONSIVE MUTATION BASED ON CLINICAL AND/OR IN VITRO DATA COMPRISING ADMINISTERING A COMPOSITION ACCORDING TO CLAIM 1 OF US 11147770
- U-4061 TREATMENT OF CF IN PATIENTS AGED 2 TO <6 YEARS WHO HAVE IN THE CFTR GENE AT LEAST ONE F508DEL MUTATION OR A RESPONSIVE MUTATION BASED ON CLINICAL AND/OR IN VITRO DATA WITH ELEXACAFTOR, TEZACAFTOR, AND IVACAFTOR
- U-4062 TREATMENT OF CF IN PATIENTS AGED 2 TO <6 YEARS WHO HAVE IN THE CFTR GENE AT LEAST ONE F508DEL MUTATION OR A RESPONSIVE MUTATION BASED ON CLINICAL AND/OR IN VITRO DATA COMPRISING ADMINISTERING A COMPOSITION ACCORDING TO CLAIM 1 OF US 10272046
- U-4063 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 6 YEARS AND OLDER WHO HAVE IN THE CFTR GENE AT LEAST ONE F508DEL MUTATION OR A RESPONSIVE MUTATION BASED ON CLINICAL AND/OR IN VITRO DATA BY ADMINISTERING THE COMPOSITION RECITED IN US 11564916 CLAIM 1
- U-4064 TREATMENT OF CF IN PATIENTS AGED 2 TO <6 YEARS WHO HAVE IN THE CFTR GENE AT LEAST ONE F508DEL MUTATION OR A RESPONSIVE MUTATION BASED ON CLINICAL AND/OR IN VITRO DATA WITH AN EFFECTIVE AMOUNT OF ELX, TEZ, AND IVA
- U-4065 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 2 TO <6 YEARS WHO HAVE IN THE CFTR GENE AT LEAST ONE F508DEL MUTATION OR A RESPONSIVE MUTATION BASED ON CLINICAL AND/OR IN VITRO DATA WITH A COMPOSITION ACCORDING TO CLAIM 1 OF US 10081621
- U-4066 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 2 TO <6 YEARS WHO HAVE IN THE CFTR GENE AT LEAST ONE F508DEL MUTATION OR A RESPONSIVE MUTATION BASED ON CLINICAL AND/OR IN VITRO DATA WITH ELEXACAFTOR, TEZACAFTOR, AND IVACAFTOR
- U-4067 TREATMENT OF CF IN PATIENTS AGED 6 YEARS AND OLDER WHO HAVE IN THE CFTR GENE AT LEAST ONE F508DEL MUTATION OR A RESPONSIVE MUTATION BASED ON CLINICAL AND/OR IN VITRO DATA BY ADMINISTERING DAILY ELX (200 MG); TEZ; AND IVA
- U-4068 TREATMENT OF CF IN PATIENTS AGED 2 TO <6 YEARS HAVING IN THE CFTR GENE AT LEAST ONE F508DEL MUTATION OR A RESPONSIVE MUTATION BASED ON CLINICAL AND/OR IN VITRO DATA USING A SOLID COMPOSITION OF ELX, TEZ, AMORPHOUS IVA, AND < ~30% CRYSTALLINE IVA

PATENT AND EXCLUSIVITY TERMS

ADB 223 of 225

PATENT USE

- U-4069 TREATMENT OF CF IN PATIENTS AGED 6 YEARS AND OLDER WHO HAVE IN THE CFTR GENE AT LEAST ONE F508DEL MUTATION OR A RESPONSIVE MUTATION BASED ON CLINICAL AND/OR IN VITRO DATA BY ADMINISTERING DAILY ELX (100 MG); TEZ; AND IVA
- U-4070 TREATMENT OF CF IN PATIENTS AGED 6 YEARS AND OLDER WHO HAVE IN THE CFTR GENE AT LEAST ONE F508DEL MUTATION OR A RESPONSIVE MUTATION BASED ON CLINICAL AND/OR IN VITRO DATA WITH A COMPOUND OF CLAIM 1 OR COMPOSITION OF CLAIM 29 OF US11426407
- U-4071 TREATMENT OF CF IN PATIENTS AGED 6 YEARS AND OLDER WHO HAVE IN THE CFTR GENE AT LEAST ONE F508DEL MUTATION OR A RESPONSIVE MUTATION BASED ON CLINICAL AND/OR IN VITRO DATA WITH A COMPOSITION ACCORDING TO CLAIM 1 OF US11179367
- U-4072 TREATMENT OF A MODERATE TO MILD CLINICAL PHENOTYPE OF CF IN PATIENTS AGED 2 TO <6 YEARS WHO HAVE IN THE CFTR GENE AT LEAST ONE F508DEL MUTATION OR A RESPONSIVE MUTATION BASED ON CLINICAL AND/OR IN VITRO DATA WITH ELX, TEZ, AND IVA
- U-4073 TREATMENT OF CF IN PATIENTS AGED 6 YEARS AND OLDER WHO HAVE IN THE CFTR GENE AT LEAST ONE F508DEL MUTATION OR A RESPONSIVE MUTATION BASED ON CLINICAL AND/OR IN VITRO DATA WITH ELEXACAFTOR, TEZACAFTOR, AND IVACAFTOR
- U-4074 TREATMENT OF CF IN PATIENTS AGED 6 YEARS AND OLDER WHO HAVE IN THE CFTR GENE AT LEAST ONE F508DEL MUTATION OR A RESPONSIVE MUTATION BASED ON CLINICAL AND/OR IN VITRO DATA WITH AN EFFECTIVE AMOUNT OF ELX, TEZ, AND IVA
- U-4075 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 6 YEARS AND OLDER WHO HAVE IN THE CFTR GENE AT LEAST ONE F508DEL MUTATION OR A RESPONSIVE MUTATION BASED ON CLINICAL AND/OR IN VITRO DATA WITH A COMPOSITION ACCORDING TO CLAIM 1 OF US 10081621
- U-4076 TREATMENT OF CF IN PATIENTS AGED 2 TO <6 YEARS WHO HAVE IN THE CFTR GENE AT LEAST ONE F508DEL MUTATION GENE OR A RESPONSIVE MUTATION BASED ON CLINICAL AND/OR IN VITRO DATA WITH ELEXACAFTOR, TEZACAFTOR, AND IVACAFTOR
- U-4077 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 6 YEARS AND OLDER WHO HAVE IN THE CFTR GENE AT LEAST ONE F508DEL MUTATION OR A RESPONSIVE MUTATION BASED ON CLINICAL AND/OR IN VITRO DATA WITH ELEXACAFTOR, TEZACAFTOR, AND IVACAFTOR
- U-4078 TREATMENT OF CF IN PATIENTS AGED 6 YEARS AND OLDER HAVING IN THE CFTR GENE AT LEAST ONE F508DEL MUTATION OR A RESPONSIVE MUTATION BASED ON CLINICAL AND/OR IN VITRO DATA USING A SOLID COMPOSITION OF ELX, TEZ, AMORPHOUS IVA, AND < ~30% CRYSTALLINE IVA
- U-4079 TREATMENT OF A MODERATE TO MILD CLINICAL PHENOTYPE OF CF IN PATIENTS AGED 6 YEARS AND OLDER WHO HAVE IN THE CFTR GENE AT LEAST ONE F508DEL MUTATION OR A RESPONSIVE MUTATION BASED ON CLINICAL AND/OR IN VITRO DATA WITH ELX, TEZ, AND IVA
- U-4080 TREATMENT OF CF IN PATIENTS AGED 6 YEARS AND OLDER WHO HAVE AT LEAST ONE F508DEL MUTATION OR ANOTHER RESPONSIVE MUTATION IN THE CFTR GENE WITH A SOLID PHARMACEUTICAL COMPOSITION COMPRISING VNZ, TEZ, AMORPHOUS D-IVA, AND <30% CRYSTALLINE D-IVA
- U-4081 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 6 YEARS AND OLDER WHO HAVE AT LEAST ONE F508DEL MUTATION IN THE CFTR GENE WITH AN EFFECTIVE AMOUNT OF A PHARMACEUTICAL COMPOSITION COMPRISING VANZACAFTOR, TEZACAFTOR, AND DEUTIVACAFTOR
- U-4082 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 6 YEARS AND OLDER WHO HAVE AT LEAST ONE F508DEL MUTATION OR ANOTHER RESPONSIVE MUTATION IN THE CFTR GENE WITH VANZACAFTOR, TEZACAFTOR, AND DEUTIVACAFTOR
- U-4083 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 6 YEARS AND OLDER WHO HAVE AT LEAST ONE F508DEL MUTATION OR ANOTHER RESPONSIVE MUTATION IN THE CFTR GENE WITH VANZACAFTOR CALCIUM SALT HYDRATE FORM D, TEZACAFTOR, AND DEUTIVACAFTOR
- U-4084 TREATMENT OF A MODERATE TO MILD CLINICAL PHENOTYPE OF CF IN PATIENTS AGED 6 YEARS AND OLDER WHO HAVE AT LEAST ONE F508DEL MUTATION OR ANOTHER RESPONSIVE MUTATION IN THE CFTR GENE WITH A PHARMACEUTICAL COMPOSITION COMPRISING VNZ, TEZ, AND D-IVA
- U-4085 REDUCTION IN LOSS OF KIDNEY FUNCTION IN ADULTS WITH PRIMARY IMMUNOGLOBULIN A NEPHROPATHY (IGAN) WHO ARE AT RISK OF DISEASE PROGRESSION, BY RELEASE OF BUDESONIDE FROM THE FORMULATION
- U-4086 REDUCTION OF PROTEINURIA IN ADULTS WITH PRIMARY IMMUNOGLOBULIN A NEPHROPATHY (IGAN) WHO ARE AT RISK OF DISEASE PROGRESSION, BY RELEASE OF BUDESONIDE FROM THE FORMULATION
- U-4087 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 6 YEARS AND OLDER WHO HAVE AT LEAST ONE F508DEL MUTATION OR ANOTHER RESPONSIVE MUTATION IN THE CFTR GENE WITH

PATENT AND EXCLUSIVITY TERMS

ADB 224 of 225

PATENT USE

- VNZ, D-IVA, AND A COMPOSITION ACCORDING TO CLAIM 1 OF US10081621
- U-4088 TREATMENT OF CF IN PATIENTS AGED 6 YEARS AND OLDER WHO HAVE AT LEAST ONE F508DEL MUTATION OR ANOTHER RESPONSIVE MUTATION IN THE CFTR GENE WITH 21.4 MG OF VNZ CALCIUM SALT HYDRATE FORM D, 100 MG OF TEZ, AND 250 MG OF D-IVA DAILY
- U-4089 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 6 YEARS AND OLDER WHO HAVE AT LEAST ONE F508DEL MUTATION OR ANOTHER RESPONSIVE MUTATION IN THE CFTR GENE WITH VNZ AND A COMPOSITION ACCORDING TO CLAIM 1 OF US11951212
- U-4090 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 6 YEARS AND OLDER WHO HAVE AT LEAST ONE F508DEL MUTATION OR ANOTHER RESPONSIVE MUTATION IN THE CFTR GENE WITH AN EFFECTIVE AMOUNT OF VANZACAFOTOR, TEZACAFOTOR, AND DEUTIVACAFOTOR
- U-4091 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 6 YEARS AND OLDER WHO HAVE AN R117H MUTATION IN THE CFTR GENE WITH VANZACAFOTOR, TEZACAFOTOR, AND DEUTIVACAFOTOR
- U-4092 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 6 YEARS AND OLDER WHO HAVE AT LEAST ONE F508DEL MUTATION OR ANOTHER RESPONSIVE MUTATION IN THE CFTR GENE WITH VNZ AND A COMPOSITION ACCORDING TO CLAIM 1 OF US10206877
- U-4093 TREATMENT OF CF IN PATIENTS 6 YEARS AND OLDER WITH AT LEAST ONE F508DEL OR G551D AND AN A455E, 2789+5G->A, OR 3849+10KBC->T CFTR MUTATION, COMPRISING CONCURRENT COADMINISTRATION OF THE COMPOSITIONS OF CLAIM 1 OF US 10058546 FURTHER COMPRISING VNZ
- U-4094 TREATMENT OF CF IN PATIENTS AGED 6 YEARS AND OLDER WHO HAVE AT LEAST ONE F508DEL MUTATION OR ANOTHER RESPONSIVE MUTATION IN THE CFTR GENE WITH A COMPOSITION COMPRISING VNZ, TEZ, AND D-IVA WITH AT LEAST 90% ISOTOPIC ENRICHMENT FOR EACH DEUTERIUM ATOM
- U-4095 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 6 YEARS AND OLDER WHO HAVE AT LEAST ONE F508DEL MUTATION OR ANOTHER RESPONSIVE MUTATION IN THE CFTR GENE WITH A COMPOSITION ACCORDING TO CLAIM 1 OF US11564916, VNZ, AND TEZ
- U-4096 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 6 YEARS AND OLDER WHO HAVE AT LEAST ONE F508DEL MUTATION OR ANOTHER RESPONSIVE MUTATION IN THE CFTR GENE WITH VANZACAFOTOR, DEUTIVACAFOTOR, AND A SOLID DISPERSION ACCORDING TO CLAIM 1 OF US 11,578,062
- U-4097 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 6 YEARS AND OLDER WHO HAVE AT LEAST ONE F508DEL MUTATION IN THE CFTR GENE, COMPRISING CONCURRENT COADMINISTRATION OF THE COMPOSITIONS OF CLAIM 1 OF US 9,012,496 FURTHER COMPRISING VNZ
- U-4098 TREATMENT OF CF IN PATIENTS AGED 6 YEARS AND OLDER WHO HAVE AT LEAST ONE F508DEL MUTATION OR ANOTHER RESPONSIVE MUTATION IN THE CFTR GENE WITH AN EFFECTIVE AMOUNT OF VNZ, TEZ, AND A COMPOSITION ACCORDING TO CLAIM 3 OF US 9,512,079
- U-4099 TREATMENT OF ADULT PATIENTS WITH ANAPLASTIC LYMPHOMA KINASE (ALK)-POSITIVE LOCALLY ADVANCED OR METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) WHO HAVE NOT PREVIOUSLY RECEIVED AN ALK-INHIBITOR
- U-4100 A METHOD OF TREATING MODERATE TO SEVERE OPIOID USE DISORDER BY ADMINISTERING BUPRENORPHINE ONCE WEEKLY
- U-4101 TREATMENT OF PATIENTS WITH CASTRATION-RESISTANT PROSTATE CANCER TO WHOM RIFAMPIN IS ADMINISTERED WITH THE CO-ADMINISTRATION OF A DAILY DOSE OF 240 MG ENZALUTAMIDE
- U-4102 TREATMENT OF PATIENTS WITH NON-METASTATIC CASTRATION-SENSITIVE PROSTATE CANCER WITH BIOCHEMICAL RECURRENCE AT HIGH RISK FOR METASTASIS TO WHOM RIFAMPIN IS ADMINISTERED WITH THE CO-ADMINISTRATION OF A DAILY DOSE OF 240 MG ENZALUTAMIDE
- U-4103 TREATMENT OF PATIENTS WITH METASTATIC CASTRATION-SENSITIVE PROSTATE CANCER TO WHOM RIFAMPIN IS ADMINISTERED WITH THE CO-ADMINISTRATION OF A DAILY DOSE OF 240 MG ENZALUTAMIDE
- U-4104 TREATMENT OF PATIENTS WITH METASTATIC CASTRATION-RESISTANT PROSTATE CANCER TO WHOM RIFAMPIN IS ADMINISTERED WITH THE CO-ADMINISTRATION OF A DAILY DOSE OF 240 MG ENZALUTAMIDE
- U-4105 USE OF VASCEPA TO LOWER TRIGLYCERIDES AND LDL-C IN AN ADULT PATIENT ON STATIN THERAPY WITH CARDIOVASCULAR-RELATED DISEASE AND ELEVATED TRIGLYCERIDE (TG) LEVELS (ABOUT 200 MG/DL TO ABOUT 500 MG/DL)
- U-4106 TREATMENT OF EXCESSIVE DAYTIME SLEEPINESS BY ADMINISTERING SOLRIAMFETOL TO A SUBJECT HAVING NO OR MILD RENAL IMPAIRMENT

PATENT AND EXCLUSIVITY TERMS