

APPROVED DRUG PRODUCTS

WITH

**THERAPEUTIC
EQUIVALENCE
EVALUATIONS**

MARCH 20, 2020 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**

2020

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 March 20, 2020.

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Therapeutic Equivalence Evaluations**

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On March 23, 2020, the Biologics Price Competition and Innovation Act of 2009 requires that an approved application for a biological product under section 505 of the Federal Food, Drug, and Cosmetic Act (FD&C Act) shall be deemed to be a license for the biological product under section 351 of the Public Health Service Act (PHS Act) (see section 7002(e)(4) of the Biologics Price Competition and Innovation Act of 2009). FDA interprets the statute to mean that an approved application for a biological product under section 505 of the FD&C Act (including an approved application for a biological product that has been discontinued from marketing, but for which FDA has not withdrawn approval of the application), will be deemed to be an approved biologics license application for that product on March 23, 2020. However, if FDA has withdrawn approval of an application for a biological product under section 505 of the FD&C Act, the application will be removed from the Orange Book on March 23, 2020, and will not transition. As a result, on March 23, 2020, FDA removed from the Orange Book the listings for "biological products" that have been approved in applications under section 505 of the FD&C Act, based on the Agency's position that these products are no longer "listed drugs." For additional information, see FDA's guidance on [Interpretation of the "Deemed to be a License" Provision of the Biologics Price Competition and Innovation Act of 2009](#) (December 2018) and FDA's guidance on [The "Deemed to Be a License" Provision of the BPCI Act Questions and Answers](#).

To enhance transparency, we are making available an archival copy of the Orange Book as of March 20, 2020, the last working day before the March 23, 2020 transition date. This archival copy contains information available in the Orange Book as of the morning of March 20, 2020, and generally includes information, such as patent and exclusivity information, for "biological products" that have been approved under section 505 of the FD&C Act as of March 19, 2020.

PREFACE TO MARCH 20, 2020 EDITION

1.0 INTRODUCTION

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 efficacy 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 health care 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 corrections and additions. 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 abbreviated new drug application (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 Legal and Regulatory Support, 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.

1.1 Content and Exclusion

The Orange Book is composed of four parts: (1) approved prescription drug products with therapeutic equivalence evaluations; (2) approved over-the-counter (OTC) drug products for those drugs that may not be marketed without NDAs or ANDAs because they are not covered under existing OTC monographs; (3) drug products with approval under Section 505 of the FD&C Act administered by the Center for Biologics Evaluation and Research; 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 efficacy reasons subsequent to being discontinued from marketing.¹ This publication also includes indices of prescription and OTC drug products by proprietary name (brand name or trade name) or, if no proprietary name exists, established name of the active ingredient and by applicant name, which have been abbreviated for this publication. Established names for active ingredients generally conform to compendial names or *United States Adopted Names* (USAN) as described in 21 CFR 299.4(e). A list of uniform terms is provided in Appendix C.

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

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

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 efficacy and for which new drug applications 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. The Agency will list the drug product in the Orange Book and the date of approval as determined under Section 505(x).

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, 250mg capsules vs. tetracycline phosphate complex, 250mg capsules; quinidine sulfate, 200mg tablets vs. quinidine sulfate, 200mg 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.

² 21 CFR 314.3(b).

³ See 21 CFR 314.3(b).

⁴ 21 CFR 314.3(b).

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 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 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

⁵ 21 CFR 314.3(b).

⁶ 21 CFR 314.3(b).

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. 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 200MG 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 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.⁷

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.⁸ Section 505(j)(8)(B) of the FD&C Act describes certain conditions under which a test drug and reference listed drug (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,

⁷ 21 CFR 314.3(b).

⁸ 21 CFR 314.3(b).

bioequivalence may sometimes be demonstrated through comparative clinical trials or pharmacodynamic studies.⁹

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.¹⁰

1.4 Reference Listed Drug and Reference Standard

A reference listed drug is the listed drug¹¹ identified by FDA as the drug product upon which an applicant relies in seeking approval of its ANDA.¹² Generally, a reference listed drug is a drug product approved in a new drug application under Section 505(c) of the FD&C Act based on full reports of investigations of safety and effectiveness. For an ANDA based on an approved suitability petition (a petitioned ANDA), the reference listed drug generally is the listed drug referenced in the approved suitability petition.¹³

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.¹⁴ FDA generally selects a single reference standard that ANDA applicants must use in *in vivo* bioequivalence testing. Ordinarily, FDA will select the reference listed drug as the reference standard. However, in some instances, the reference listed drug and the reference standard may be different. For example, where the reference listed drug has been withdrawn from sale FDA may select an ANDA as the reference standard.

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

In some instances when FDA has not designated a listed drug as a reference listed drug, such listed drug may be shielded from generic competition. If FDA has not designated a reference listed drug 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

⁹ 21 CFR 320.24.

¹⁰ 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 d/////ocuments 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>.

¹¹ 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" (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)).

¹² 21 CFR 314.3(b).

¹³ 21 CFR 314.94(a)(3)(i).

¹⁴ 21 CFR 314.3(b).

designate a reference listed drug 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 drug products listed under the same heading with two or more reference listed drugs.

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

If an applicant has a question related to the appropriate reference standard, it is recommended that an applicant planning to conduct an *in vivo* bioequivalence study 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 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 Directory¹⁵ 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

¹⁵ <https://www.fda.gov/drugs/drug-approvals-and-databases/national-drug-code-directory>.

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 efficacy 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, 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 product (e.g., a particular strength of an approved drug) as therapeutically equivalent to other pharmaceutically equivalent 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 to the reference product 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 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 product 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 product. When such changes in the listed drug product 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, 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 health care professionals from converting pharmaceutically different concentrations into pharmaceutical equivalents using accepted professional practice.

Where package size variations have therapeutic implications, products so packaged have not been considered pharmaceutically equivalent. For example, some oral contraceptives are supplied in 21-tablet and 28-tablet packets; the 28-tablet packets contain 7 placebo or iron tablets. These two packaging configurations are not regarded as pharmaceutically equivalent; thus, they are not designated as therapeutically equivalent.

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 reference listed drug of the same strength has been designated under the same heading. If a study is submitted that demonstrates bioequivalence to a reference listed drug product, the generic product will be given the same three-character code as the reference listed drug 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 available as dry powders for reconstitution, concentrated sterile solutions for dilution, or sterile solutions ready for injection are pharmaceutical alternative drug products. 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 similarly labeled. 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.

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 parenteral drug products generally is identified by both the total drug content and the concentration of drug substance in a container approved by FDA.¹⁶ In the past, the strength of liquid parenteral drug products in the Orange Book has not been fully displayed. Rather, the strength of liquid parenteral drug products in the Orange Book has been displayed in terms of concentration, expressed as x mg/mL. 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 displays the strength of all 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. 50 mg/mL. For example, if this application had a 20 mL and 60 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. 1 gm/20 mL (50 mg/mL) and 3 gm/60 mL (50 mg/mL).

AT Topical products

There are a variety of topical dosage forms available for dermatologic, ophthalmic, otic, rectal, and vaginal administration, including creams,

¹⁶ The strengths of certain parenteral drug products, including contrast agents, may be expressed as a percentage.

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.

"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 are 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 ingredient meeting these criteria as having a potential bioequivalence problem even in the absence of positive data demonstrating inequivalence. Pharmaceutically equivalent products containing these 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 reference listed drug 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.¹⁷ Because there are multiple reference listed drugs for levothyroxine sodium tablets and some reference listed drugs' sponsors have conducted studies to establish their drugs' therapeutic equivalence to other reference listed drugs, 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 a reference listed drug or reference standard, it may be difficult to determine to which therapeutic equivalence code the reference listed drugs 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 reference listed drug and reference standard, but it is unclear that the reference listed drug and reference standard designations are associated with the AB1 therapeutic equivalence code.

Accordingly, FDA provides the following chart, which identifies (1) a reference listed drug for each therapeutic equivalence code in the Orange Book and (2) and the reference standard products in the Active Section of the Orange Book.¹⁸

- Therapeutic equivalence has been established between products that have the same AB+number therapeutic equivalence code (i.e. AB1, AB2, AB3 or

¹⁷ In previous editions of the Orange Book, FDA provided a chart outlining therapeutic equivalence codes for all .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.

¹⁸ 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).

- 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 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	0.3MG	N021210	RLD	RS
AB2	SYNTHROID	ABBVIE	0.3MG	N021402	RLD	RS
AB3	LEVOXYL	KING PHARMS	0.2MG	N021301	RLD	RS
AB4	THYRO-TABS	ALVOGEN GROUP HOLDINGS 4 LLC	0.3MG	N021116	RLD	-
AB4	LEVOTHYROXINE SODIUM ¹⁹	MYLAN	0.3MG	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 reference listed drug 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 reference listed drug, 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 reference listed drug 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 reference listed drug, the reference listed drug 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 reference listed drug 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

¹⁹ Lloyd's Thyro-Tabs 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 reference listed drugs (i.e., Unithroid, Synthroid, and Levoxyl) by submitting supplements that demonstrate that the generic product is bioequivalent to these other reference listed drugs 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).

to the reference listed drug that served as the basis for the approved suitability petition.²⁰ (This concept also generally applies to an ANDA applicant that utilizes a reference standard that is not a reference listed drug, as such an application must include appropriate patent certification(s) and an exclusivity statement with respect to the reference listed drug.)

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 Section of the Orange Book. If a drug product has received this exclusivity, FDA will not accept for review and/or will not approve a 505(b)(2) application or an ANDA under Section 505(j) of the FD&C Act, as applicable, in accordance with the relevant exclusivity.²¹ If the listed drug is also protected by one or more patents, the approval date for the ANDA or 505(b)(2) application 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 ANDAs and 505(b)(2) applications that might otherwise be blocked by such exclusivity. If an NDA sponsor waives its exclusivity, qualified ANDAs or 505(b)(2) applications may be accepted for review and/or approved, as applicable. An NDA for which the holder has waived its exclusivity as to all ANDAs and 505(b)(2) applications will be coded with a "W" in the Patent and Exclusivity Section of the Orange Book. The applicant whose 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,

²⁰ 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 listed drug 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).

²¹ See Patent and Exclusivity Information Addendum in the Orange Book.

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 may 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).

1.11 Discontinued Section

Those drug products in the discontinued section 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 efficacy 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 Orange Book staff 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.²² 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 the FDA Orange Book Staff 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, applicants 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 applicants 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 the Orange Book staff by the end of the month in which the product is approved to ensure that the product is not included in the "active" portions of the next published Orange Book update.

In addition, the FDA Orange Book Staff 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 the Orange Book Staff 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. An annual subscription of the PDF format may be obtained from the U.S. Government Publishing Office, <https://www.gpo.gov/>.

²² See, e.g., Section 506I(d) of the FD&C Act.

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 (dosage form, product name, strength, and application number).

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>					
THERAPEUTIC EQUIVALENCE (TE)	→	<u>AP</u>	GREENBERG PHARM	<u>25MG/ML</u>	<u>A064890</u>	001	FEB 29, 1987
CODE FOR MULTISOURCE PRODUCT	→	<u>AP</u>		<u>50MG/ML</u>	<u>A064890</u>	002	FEB 29, 1987
		<u>AP</u>		<u>75MG/ML</u>	<u>A064890</u>	003	FEB 29, 1987
		<u>AP</u>		<u>100MG/ML</u>	<u>A064890</u>	004	MAR 08, 1992
SINGLE SOURCE PRODUCT (NO TE CODE)			! 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
			! KENDRA PHARM	150MG/ML	A079444	001	OCT 31, 1999
APPLICANT	→						
AVAILABLE STRENGTH(S) OF A PRODUCT	→						
APPLICATION NUMBER AND PRODUCT 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

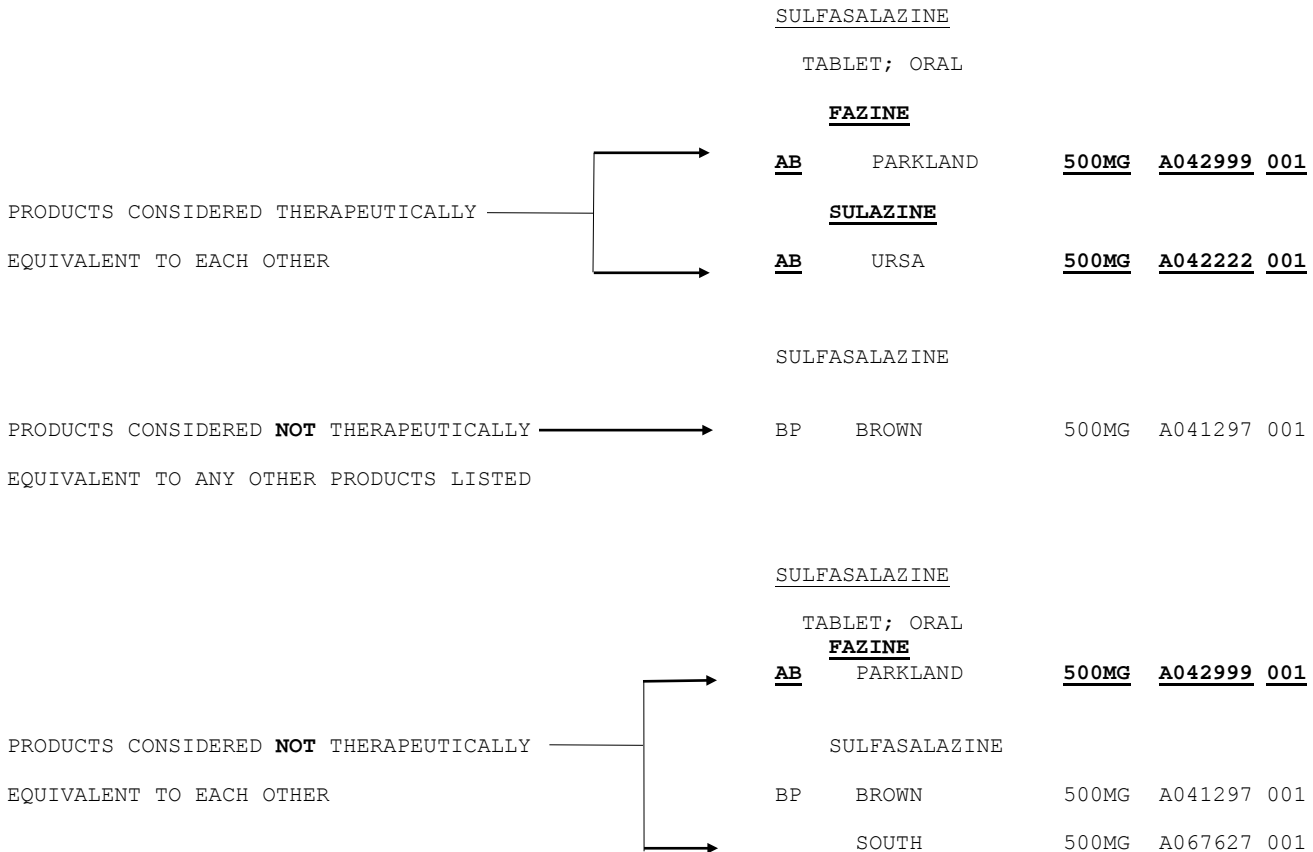
MULTIPLE INGREDIENTS WITH PRODUCT INFORMATION

ALPHABETICALLY SORTED BY		
ACTIVE INGREDIENT	→	<u>HYDRALAZINE HYDROCHLORIDE; HYDROCHLOROTHIAZIDE; RESERPINE</u>
PRODUCT INFORMATION	→	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*.



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
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TABLET;ORAL

ABACAVIR SULFATE

AB	APOTEX INC	EQ 300MG BASE	A201570 001	Dec 17, 2012
AB	AUROBINDO PHARMA LTD	EQ 300MG BASE	A077844 001	Dec 17, 2012
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
AB	STRIDES PHARMA	EQ 300MG BASE	A091050 001	Oct 28, 2016

ZIAGEN

AB	+! VIIV HLHCARE	EQ 300MG BASE	N020977 001	Dec 17, 1998
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ABACAVIR SULFATE; DOLUTEGRAVIR SODIUM; LAMIVUDINE

TABLET;ORAL

TRIUMEQ

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

TABLET;ORAL

ABACAVIR SULFATE AND LAMIVUDINE

AB	AUROBINDO PHARMA LTD	EQ 600MG BASE;300MG	A090159 001	Nov 15, 2018
AB		EQ 600MG BASE;300MG	A206151 001	Mar 28, 2017
AB	CIPLA	EQ 600MG BASE;300MG	A091144 001	Mar 28, 2017
AB	LUPIN LTD	EQ 600MG BASE;300MG	A204990 001	Mar 28, 2017
AB	TEVA PHARMS USA	EQ 600MG BASE;300MG	A079246 001	Sep 29, 2016

EPZICOM

AB	+! VIIV HLHCARE	EQ 600MG BASE;300MG	N021652 001	Aug 02, 2004
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ABACAVIR SULFATE; LAMIVUDINE; ZIDOVUDINE

TABLET;ORAL

ABACAVIR SULFATE, LAMIVUDINE AND ZIDOVUDINE

AB	LUPIN LTD	EQ 300MG BASE;150MG;300MG	A202912 001	Dec 05, 2013
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TRIZIVIR

AB	+! VIIV HLHCARE	EQ 300MG BASE;150MG;300MG	N021205 001	Nov 14, 2000
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ABALOPARATIDE

SOLUTION;SUBCUTANEOUS

TYMLOS

+!	RADIUS HEALTH INC	3.12MG/1.56ML (2MG/ML)	N208743 001	Apr 28, 2017
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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	APOTEX	250MG	A208453 001	Oct 31, 2018
AB	GLENMARK PHARMS	250MG	A209227 001	Oct 16, 2019
AB	HIKMA PHARMS	250MG	A208339 001	Oct 31, 2018
AB	MSN	250MG	A210686 001	Jul 10, 2019
AB	MYLAN	250MG	A208446 001	Oct 31, 2018
AB	QILU	250MG	A212462 001	Sep 27, 2019
AB	RISING	250MG	A208371 001	Feb 25, 2019
AB	TEVA PHARMS USA	250MG	A208432 001	Oct 31, 2018
AB	WOCKHARDT BIO AG	250MG	A208380 001	Feb 27, 2019

ZYTIGA

AB	+ JANSSEN BIOTECH	250MG	N202379 001	Apr 28, 2011
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YONSA

+!	SUN PHARMA GLOBAL	125MG	N210308 001	May 22, 2018
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PRESCRIPTION DRUG PRODUCT LIST

ABIRATERONE ACETATE

TABLET; ORAL

ZYTIGA

+! JANSSEN BIOTECH 500MG N202379 002 Apr 14, 2017

ACALABRUTINIB

CAPSULE; ORAL

CALQUENCE

+! ASTRAZENECA 100MG N210259 001 Oct 31, 2017

ACAMPROSATE CALCIUM

TABLET, DELAYED RELEASE; ORAL

ACAMPROSATE CALCIUM

AB	!	GLENMARK GENERICS	333MG	A202229	001	Jul 16, 2013
AB		MYLAN	333MG	A200142	001	Mar 11, 2014
AB		ZYDUS PHARMS	333MG	A205995	001	May 26, 2017

ACARBOSE

TABLET; ORAL

ACARBOSE

AB		EMCURE PHARMS LTD	25MG	A202271	001	Feb 07, 2012
AB			50MG	A202271	002	Feb 07, 2012
AB			100MG	A202271	003	Feb 07, 2012
AB		HIKMA	25MG	A078470	001	May 07, 2008
AB			50MG	A078470	002	May 07, 2008
AB			100MG	A078470	003	May 07, 2008
AB		IMPAX LABS	25MG	A078441	001	May 14, 2009
AB			50MG	A078441	002	May 14, 2009
AB			100MG	A078441	003	May 14, 2009
AB	!	STRIDES PHARMA	25MG	A090912	001	Jul 27, 2011
AB			50MG	A090912	002	Jul 27, 2011
AB			100MG	A090912	003	Jul 27, 2011
AB		VIRTUS PHARM	25MG	A091343	001	Oct 17, 2013
AB			50MG	A091343	002	Oct 17, 2013
AB			100MG	A091343	003	Oct 17, 2013
AB		WATSON LABS	25MG	A077532	001	May 07, 2008
AB			50MG	A077532	002	May 07, 2008
AB			100MG	A077532	003	May 07, 2008

ACEBUTOLOL HYDROCHLORIDE

CAPSULE; ORAL

ACEBUTOLOL HYDROCHLORIDE

AB	!	AMNEAL PHARM	EQ 200MG BASE	A075047	001	Dec 30, 1999
AB	!		EQ 400MG BASE	A075047	002	Dec 30, 1999
AB		MYLAN	EQ 200MG BASE	A074288	001	Apr 24, 1995
AB			EQ 400MG BASE	A074288	002	Apr 24, 1995

ACETAMINOPHEN

SOLUTION; INTRAVENOUS

ACETAMINOPHEN

AP		CUSTOPHARM INC	1GM/100ML (10MG/ML)	A202605	001	Jun 13, 2016
AP		SANDOZ INC	1GM/100ML (10MG/ML)	A204052	001	Mar 22, 2016
OFIRMEV						
AP	+	MALLINCKRODT HOSP	1GM/100ML (10MG/ML)	N022450	001	Nov 02, 2010
ACETAMINOPHEN						
		FRESENIUS KABI USA	1GM/100ML (10MG/ML)	N204767	001	Oct 28, 2015

ACETAMINOPHEN; BENZHYDROCODONE HYDROCHLORIDE

TABLET; ORAL

APADAZ

+	KVK TECH INC	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

BUTALBITAL AND ACETAMINOPHEN

AA		GRANULES PHARMS	300MG;50MG	A213115	001	Nov 22, 2019
AA	!	MAYNE PHARMA INC	300MG;50MG	A207313	001	Dec 27, 2017

TABLET; ORAL

BUTALBITAL AND ACETAMINOPHEN

AA		CNTY LINE PHARMS	300MG;50MG	A207635	001	Jun 05, 2017
AA			325MG;50MG	A205120	001	Oct 30, 2015
AA		LARKEN LABS INC	325MG;50MG	A203484	002	Dec 04, 2015
AA		MIKART	300MG;50MG	A207386	001	Nov 15, 2016
AA	!	NEXGEN PHARMA	300MG;50MG	A090956	001	Aug 23, 2011

PRESCRIPTION DRUG PRODUCT LISTACETAMINOPHEN; BUTALBITAL

TABLET;ORAL

BUTAPAP

AA	!	MIKART	<u>325MG;50MG</u>	<u>A089987</u>	<u>001</u>	Oct 26, 1992
		ALLZITAL				
	!	LARKEN LABS INC	325MG;25MG	A203484	001	Dec 04, 2015

ACETAMINOPHEN; BUTALBITAL; CAFFEINE

CAPSULE;ORAL

BUTALBITAL, ACETAMINOPHEN AND CAFFEINE

AA		AUROLIFE PHARMA LLC	<u>325MG;50MG;40MG</u>	<u>A204733</u>	<u>001</u>	Sep 26, 2018
AA		LANNETT CO INC	<u>300MG;50MG;40MG</u>	<u>A212082</u>	<u>001</u>	Dec 17, 2019
AA			<u>325MG;50MG;40MG</u>	<u>A212083</u>	<u>001</u>	Dec 17, 2019
AA		MAYNE PHARMA INC	<u>300MG;50MG;40MG</u>	<u>A210817</u>	<u>001</u>	Dec 17, 2019
AA	!		<u>325MG;50MG;40MG</u>	<u>A089007</u>	<u>001</u>	Mar 17, 1986
AA	!	NEXGEN PHARMA	<u>300MG;50MG;40MG</u>	<u>A040885</u>	<u>001</u>	Nov 16, 2009
AA		NUVO PHARMS INC	<u>300MG;50MG;40MG</u>	<u>A207118</u>	<u>001</u>	Oct 28, 2016
AA		WRASER PHARMS LLC	<u>300MG;50MG;40MG</u>	<u>A206615</u>	<u>001</u>	Aug 04, 2017

SOLUTION;ORAL

BUTALBITAL, ACETAMINOPHEN AND CAFFEINE

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

BUTALBITAL, ACETAMINOPHEN AND CAFFEINE

AA		ABHAI LLC	<u>325MG;50MG;40MG</u>	<u>A211106</u>	<u>001</u>	Sep 26, 2018
AA		ACTAVIS LABS UT INC	<u>325MG;50MG;40MG</u>	<u>A088616</u>	<u>001</u>	Nov 09, 1984
AA		CNTY LINE PHARMS	<u>325MG;50MG;40MG</u>	<u>A204984</u>	<u>001</u>	Jan 10, 2017
AA		LANNETT CO INC	<u>325MG;50MG;40MG</u>	<u>A200243</u>	<u>001</u>	Sep 13, 2012
AA		MIKART	<u>325MG;50MG;40MG</u>	<u>A089175</u>	<u>001</u>	Jan 21, 1987
AA		NEXGEN PHARMA INC	<u>325MG;50MG;40MG</u>	<u>A209587</u>	<u>001</u>	Oct 31, 2018
AA		SPECGX LLC	<u>325MG;50MG;40MG</u>	<u>A087804</u>	<u>001</u>	Jan 24, 1985
AA	!	VINTAGE PHARMS	<u>325MG;50MG;40MG</u>	<u>A040511</u>	<u>001</u>	Aug 27, 2003

ACETAMINOPHEN; BUTALBITAL; CAFFEINE; CODEINE PHOSPHATE

CAPSULE;ORAL

BUTALBITAL, ACETAMINOPHEN, CAFFEINE AND CODEINE PHOSPHATE

AB		NEXGEN PHARMA INC	<u>325MG;50MG;40MG;30MG</u>	<u>A076560</u>	<u>001</u>	Jun 10, 2004
AB		VINTAGE PHARMS	<u>325MG;50MG;40MG;30MG</u>	<u>A075929</u>	<u>001</u>	Apr 22, 2002
		<u>FIORICET W/ CODEINE</u>				
AB	+	ACTAVIS LABS UT INC	<u>325MG;50MG;40MG;30MG</u>	<u>N020232</u>	<u>001</u>	Jul 30, 1992
		BUTALBITAL, ACETAMINOPHEN, CAFFEINE AND CODEINE PHOSPHATE				
		NEXGEN PHARMA INC	300MG;50MG;40MG;30MG	A076560	002	Jul 19, 2012

ACETAMINOPHEN; CAFFEINE; DIHYDROCODEINE BITARTRATE

CAPSULE;ORAL

TREZIX

		WRASER PHARMS LLC	320.5MG;30MG;16MG	A204785	001	Nov 26, 2014
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TABLET;ORAL

ACETAMINOPHEN, CAFFEINE AND DIHYDROCODEINE BITARTRATE

		LARKEN LABS INC	325MG;30MG;16MG	A204209	001	Sep 30, 2016
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ACETAMINOPHEN; CODEINE PHOSPHATE

SOLUTION;ORAL

ACETAMINOPHEN AND CODEINE PHOSPHATE

AA		ANDA REPOSITORY	<u>120MG/5ML;12MG/5ML</u>	<u>A089450</u>	<u>001</u>	Oct 27, 1992
AA		HI TECH PHARMA	<u>120MG/5ML;12MG/5ML</u>	<u>A040119</u>	<u>001</u>	Apr 26, 1996
AA	!	PHARM ASSOC	<u>120MG/5ML;12MG/5ML</u>	<u>A087508</u>	<u>001</u>	
AA		WOCKHARDT BIO AG	<u>120MG/5ML;12MG/5ML</u>	<u>A087006</u>	<u>001</u>	

TABLET;ORAL

ACETAMINOPHEN AND CODEINE PHOSPHATE

AA		AMNEAL PHARMS NY	<u>300MG;30MG</u>	<u>A040779</u>	<u>001</u>	May 29, 2008
AA		AUROLIFE PHARMA LLC	<u>300MG;15MG</u>	<u>A202800</u>	<u>001</u>	Apr 15, 2013
AA			<u>300MG;30MG</u>	<u>A202800</u>	<u>002</u>	Apr 15, 2013
AA			<u>300MG;60MG</u>	<u>A202800</u>	<u>003</u>	Apr 15, 2013
AA		ELITE LABS INC	<u>300MG;15MG</u>	<u>A212418</u>	<u>001</u>	Sep 10, 2019
AA			<u>300MG;30MG</u>	<u>A212418</u>	<u>002</u>	Sep 10, 2019
AA			<u>300MG;60MG</u>	<u>A212418</u>	<u>003</u>	Sep 10, 2019
AA		EYWA	<u>300MG;15MG</u>	<u>A211610</u>	<u>001</u>	Jun 27, 2019
AA			<u>300MG;30MG</u>	<u>A211610</u>	<u>002</u>	Jun 27, 2019
AA			<u>300MG;60MG</u>	<u>A211610</u>	<u>003</u>	Jun 27, 2019
AA	!	SPECGX LLC	<u>300MG;15MG</u>	<u>A040419</u>	<u>001</u>	May 31, 2001
AA			<u>300MG;30MG</u>	<u>A040419</u>	<u>002</u>	May 31, 2001
AA			<u>300MG;60MG</u>	<u>A040419</u>	<u>003</u>	May 31, 2001
AA		SUN PHARM INDS LTD	<u>300MG;30MG</u>	<u>A085868</u>	<u>001</u>	
AA			<u>300MG;60MG</u>	<u>A087083</u>	<u>001</u>	

PRESCRIPTION DRUG PRODUCT LIST

ACETAMINOPHEN; CODEINE PHOSPHATE

TABLET; ORAL

ACETAMINOPHEN AND CODEINE PHOSPHATE

<u>AA</u>	TEVA	<u>300MG;15MG</u>	<u>A088627</u>	<u>001</u>	Mar 06, 1985
<u>AA</u>		<u>300MG;30MG</u>	<u>A088628</u>	<u>001</u>	Mar 06, 1985
<u>AA</u>	!	<u>300MG;60MG</u>	<u>A088629</u>	<u>001</u>	Mar 06, 1985
<u>AA</u>	VINTAGE	<u>300MG;15MG</u>	<u>A089990</u>	<u>001</u>	Sep 30, 1988
<u>AA</u>		<u>300MG;30MG</u>	<u>A089805</u>	<u>001</u>	Sep 30, 1988
<u>AA</u>	VINTAGE PHARMS	<u>300MG;60MG</u>	<u>A089828</u>	<u>001</u>	Sep 30, 1988
<u>TYLENOL W/ CODEINE NO. 3</u>					
<u>AA</u>	!	JANSSEN PHARMS	<u>300MG;30MG</u>	<u>A085055</u>	<u>003</u>
<u>TYLENOL W/ CODEINE NO. 4</u>					
<u>AA</u>		JANSSEN PHARMS	<u>300MG;60MG</u>	<u>A085055</u>	<u>004</u>

ACETAMINOPHEN; HYDROCODONE BITARTRATE

SOLUTION; ORAL

HYDROCODONE BITARTRATE AND ACETAMINOPHEN

<u>AA</u>	!	ANDA REPOSITORY	<u>325MG/15ML;7.5MG/15ML</u>	<u>A040482</u>	<u>001</u>	Sep 25, 2003
<u>AA</u>		GENUS	<u>325MG/15ML;7.5MG/15ML</u>	<u>A040894</u>	<u>001</u>	Jul 19, 2011
<u>AA</u>		PHARM ASSOC	<u>325MG/15ML;7.5MG/15ML</u>	<u>A040838</u>	<u>001</u>	May 10, 2013
<u>AA</u>		VISTAPHARM	<u>325MG/15ML;7.5MG/15ML</u>	<u>A200343</u>	<u>001</u>	Jan 25, 2012
<u>AA</u>		WES PHARMA INC	<u>325MG/15ML;7.5MG/15ML</u>	<u>A211023</u>	<u>001</u>	Mar 08, 2019
	!	MIKART	300MG/15ML;10MG/15ML	A040881	001	Feb 25, 2010
	!	PHARM ASSOC	325MG/15ML;10MG/15ML	A040834	001	Apr 18, 2008

TABLET; ORAL

ANEXSIA 5/325

<u>AA</u>		SPECGX LLC	<u>325MG;5MG</u>	<u>A040409</u>	<u>001</u>	Oct 20, 2000
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ANEXSIA 7.5/325

<u>AA</u>		SPECGX LLC	<u>325MG;7.5MG</u>	<u>A040405</u>	<u>001</u>	Sep 08, 2000
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HYDROCODONE BITARTRATE AND ACETAMINOPHEN

<u>AA</u>		ABHAI LLC	<u>300MG;5MG</u>	<u>A209036</u>	<u>001</u>	Jun 21, 2017
<u>AA</u>			<u>300MG;7.5MG</u>	<u>A209036</u>	<u>002</u>	Jun 21, 2017
<u>AA</u>			<u>300MG;10MG</u>	<u>A209036</u>	<u>003</u>	Jun 21, 2017
<u>AA</u>			<u>325MG;5MG</u>	<u>A209037</u>	<u>001</u>	Jun 21, 2017
<u>AA</u>			<u>325MG;7.5MG</u>	<u>A209037</u>	<u>002</u>	Jun 21, 2017
<u>AA</u>			<u>325MG;10MG</u>	<u>A209037</u>	<u>003</u>	Jun 21, 2017
<u>AA</u>		AMNEAL PHARMS	<u>300MG;10MG</u>	<u>A207137</u>	<u>001</u>	Nov 29, 2016
<u>AA</u>		AMNEAL PHARMS NY	<u>300MG;5MG</u>	<u>A206869</u>	<u>001</u>	Jun 23, 2017
<u>AA</u>			<u>325MG;5MG</u>	<u>A040736</u>	<u>001</u>	Aug 25, 2006
<u>AA</u>			<u>325MG;7.5MG</u>	<u>A040746</u>	<u>002</u>	May 10, 2016
<u>AA</u>			<u>325MG;10MG</u>	<u>A040746</u>	<u>001</u>	Aug 25, 2006
<u>AA</u>	!	ANDA REPOSITORY	<u>325MG;2.5MG</u>	<u>A040846</u>	<u>001</u>	Jun 09, 2010
<u>AA</u>			<u>325MG;7.5MG</u>	<u>A040432</u>	<u>001</u>	Jan 22, 2003
<u>AA</u>		ASCENT PHARMS INC	<u>325MG;2.5MG</u>	<u>A211487</u>	<u>001</u>	Nov 07, 2018
<u>AA</u>			<u>325MG;5MG</u>	<u>A211487</u>	<u>002</u>	Nov 07, 2018
<u>AA</u>			<u>325MG;7.5MG</u>	<u>A211487</u>	<u>003</u>	Nov 07, 2018
<u>AA</u>			<u>325MG;10MG</u>	<u>A211487</u>	<u>004</u>	Nov 07, 2018
<u>AA</u>		AUROLIFE PHARMA LLC	<u>300MG;5MG</u>	<u>A207709</u>	<u>001</u>	Sep 13, 2018
<u>AA</u>			<u>300MG;7.5MG</u>	<u>A207709</u>	<u>002</u>	Sep 13, 2018
<u>AA</u>			<u>300MG;10MG</u>	<u>A207709</u>	<u>003</u>	Sep 13, 2018
<u>AA</u>			<u>325MG;5MG</u>	<u>A201013</u>	<u>001</u>	Apr 11, 2012
<u>AA</u>			<u>325MG;7.5MG</u>	<u>A201013</u>	<u>002</u>	Apr 11, 2012
<u>AA</u>			<u>325MG;10MG</u>	<u>A201013</u>	<u>003</u>	Apr 11, 2012
<u>AA</u>		EPIC PHARMA LLC	<u>325MG;5MG</u>	<u>A203863</u>	<u>001</u>	Mar 30, 2018
<u>AA</u>			<u>325MG;7.5MG</u>	<u>A203863</u>	<u>002</u>	Mar 30, 2018
<u>AA</u>			<u>325MG;10MG</u>	<u>A203863</u>	<u>003</u>	Mar 30, 2018
<u>AA</u>		GRANULES PHARMS	<u>325MG;5MG</u>	<u>A211729</u>	<u>001</u>	Jan 03, 2020
<u>AA</u>			<u>325MG;7.5MG</u>	<u>A211729</u>	<u>002</u>	Jan 03, 2020
<u>AA</u>			<u>325MG;10MG</u>	<u>A211729</u>	<u>003</u>	Jan 03, 2020
<u>AA</u>	!	MIKART	<u>300MG;5MG</u>	<u>A040658</u>	<u>001</u>	Jan 19, 2006
<u>AA</u>	!		<u>300MG;7.5MG</u>	<u>A040658</u>	<u>002</u>	Mar 24, 2006
<u>AA</u>	!		<u>300MG;10MG</u>	<u>A040658</u>	<u>003</u>	Jun 23, 2004
<u>AA</u>		NOVEL LABS INC	<u>300MG;5MG</u>	<u>A206142</u>	<u>001</u>	Nov 14, 2016
<u>AA</u>			<u>300MG;7.5MG</u>	<u>A206142</u>	<u>002</u>	Nov 14, 2016
<u>AA</u>			<u>300MG;10MG</u>	<u>A206142</u>	<u>003</u>	Nov 14, 2016
<u>AA</u>			<u>325MG;5MG</u>	<u>A206245</u>	<u>001</u>	Dec 01, 2016
<u>AA</u>			<u>325MG;7.5MG</u>	<u>A206245</u>	<u>002</u>	Dec 01, 2016
<u>AA</u>			<u>325MG;10MG</u>	<u>A206245</u>	<u>003</u>	Dec 01, 2016
<u>AA</u>		PAR PHARM	<u>300MG;5MG</u>	<u>A205001</u>	<u>001</u>	Jul 05, 2016
<u>AA</u>			<u>300MG;7.5MG</u>	<u>A205001</u>	<u>002</u>	Jul 05, 2016
<u>AA</u>			<u>300MG;10MG</u>	<u>A205001</u>	<u>003</u>	Jul 05, 2016
<u>AA</u>			<u>325MG;5MG</u>	<u>A202935</u>	<u>002</u>	Jun 15, 2016
<u>AA</u>			<u>325MG;7.5MG</u>	<u>A202935</u>	<u>003</u>	Jun 15, 2016

PRESCRIPTION DRUG PRODUCT LIST

ACETAMINOPHEN; HYDROCODONE BITARTRATE

TABLET; ORAL

HYDROCODONE BITARTRATE AND ACETAMINOPHEN

<u>AA</u>		<u>325MG;10MG</u>	<u>A202935 004</u>	Jun 15, 2016
<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>	SUN PHARM INDS INC	<u>325MG;5MG</u>	<u>A090118 001</u>	Dec 23, 2008
<u>AA</u>		<u>325MG;7.5MG</u>	<u>A090118 002</u>	Dec 23, 2008
<u>AA</u>		<u>325MG;10MG</u>	<u>A090118 003</u>	Dec 23, 2008
<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>	VINTAGE PHARMS	<u>300MG;5MG</u>	<u>A090415 001</u>	Jan 24, 2011
<u>AA</u>		<u>300MG;7.5MG</u>	<u>A090415 002</u>	Jan 24, 2011
<u>AA</u>		<u>300MG;10MG</u>	<u>A090415 003</u>	Jan 24, 2011
<u>AA</u>		<u>325MG;5MG</u>	<u>A040655 001</u>	Jan 19, 2006
<u>AA</u>		<u>325MG;7.5MG</u>	<u>A040656 001</u>	Jan 19, 2006
<u>AA</u>		<u>325MG;10MG</u>	<u>A040355 001</u>	May 31, 2000
<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
<u>AA</u>	XIROMED	<u>325MG;5MG</u>	<u>A211690 001</u>	Feb 07, 2020
<u>AA</u>		<u>325MG;7.5MG</u>	<u>A211690 002</u>	Feb 07, 2020
<u>AA</u>		<u>325MG;10MG</u>	<u>A211690 003</u>	Feb 07, 2020
<u>NORCO</u>				
<u>AA</u>	APIL	<u>325MG;2.5MG</u>	<u>A040148 004</u>	Jul 07, 2014
<u>AA</u>	!	<u>325MG;5MG</u>	<u>A040099 001</u>	Jun 25, 1997
<u>AA</u>		<u>325MG;5MG</u>	<u>A040148 005</u>	Jul 07, 2014
<u>AA</u>	!	<u>325MG;7.5MG</u>	<u>A040148 003</u>	Sep 12, 2000
<u>AA</u>	!	<u>325MG;10MG</u>	<u>A040148 001</u>	Feb 14, 1997

ACETAMINOPHEN; OXYCODONE HYDROCHLORIDE

SOLUTION; ORAL

OXYCODONE AND ACETAMINOPHEN

!	ABHAI LLC	325MG/5ML; 5MG/5ML	A211499 001	Dec 31, 2018
	MIKART INC	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
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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>	ANDA REPOSITORY	<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>	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>	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 PHARMS	<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>	LANNETT CO INC	<u>325MG; 5MG</u>	<u>A207333 001</u>	Sep 25, 2017
<u>AA</u>		<u>325MG; 10MG</u>	<u>A207333 002</u>	Sep 25, 2017
<u>AA</u>	MAYNE PHARMA INC	<u>325MG; 2.5MG</u>	<u>A090177 001</u>	Oct 20, 2008
<u>AA</u>		<u>325MG; 5MG</u>	<u>A090177 002</u>	Oct 20, 2008
<u>AA</u>		<u>325MG; 7.5MG</u>	<u>A090177 003</u>	Oct 20, 2008
<u>AA</u>		<u>325MG; 10MG</u>	<u>A090177 004</u>	Oct 20, 2008
<u>AA</u>	NESHER PHARMS	<u>325MG; 2.5MG</u>	<u>A210079 001</u>	Dec 28, 2017
<u>AA</u>		<u>325MG; 5MG</u>	<u>A210079 002</u>	Dec 28, 2017
<u>AA</u>		<u>325MG; 7.5MG</u>	<u>A210079 003</u>	Dec 28, 2017
<u>AA</u>		<u>325MG; 10MG</u>	<u>A210079 004</u>	Dec 28, 2017
<u>AA</u>	NOSTRUM 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>	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>	SUN PHARM INDS INC	<u>325MG; 2.5MG</u>	<u>A090535 001</u>	Dec 26, 2013
<u>AA</u>		<u>325MG; 5MG</u>	<u>A090535 002</u>	Dec 26, 2013
<u>AA</u>		<u>325MG; 7.5MG</u>	<u>A090535 003</u>	Dec 26, 2013
<u>AA</u>		<u>325MG; 10MG</u>	<u>A090535 004</u>	Dec 26, 2013
<u>AA</u>	VINTAGE PHARMS	<u>325MG; 2.5MG</u>	<u>A090733 001</u>	Jul 11, 2013
<u>AA</u>		<u>325MG; 5MG</u>	<u>A040105 001</u>	Jul 30, 1996
<u>AA</u>		<u>325MG; 7.5MG</u>	<u>A090734 001</u>	Jul 11, 2013
<u>AA</u>		<u>325MG; 10MG</u>	<u>A090734 002</u>	Jul 11, 2013
<u>AA</u>	WATSON LABS	<u>325MG; 5MG</u>	<u>A040171 001</u>	Oct 30, 1997
<u>AA</u>		<u>325MG; 7.5MG</u>	<u>A040535 001</u>	Sep 05, 2003
<u>AA</u>		<u>325MG; 10MG</u>	<u>A040535 002</u>	Sep 05, 2003
<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
<u>AA</u>	XIROMED	<u>325MG; 5MG</u>	<u>A207574 001</u>	Dec 13, 2016

PERCOCET

<u>AA</u>	!	VINTAGE PHARMS LLC	<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

ROXICET

<u>AA</u>		HIKMA	<u>325MG; 5MG</u>	<u>A087003 001</u>	
		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>325MG; 37.5MG</u>	<u>A090485 001</u>	Dec 09, 2009
<u>AB</u>	AUROBINDO PHARMA LTD	<u>325MG; 37.5MG</u>	<u>A207152 001</u>	Mar 22, 2017

PRESCRIPTION DRUG PRODUCT LIST

ACETAMINOPHEN; TRAMADOL HYDROCHLORIDE

TABLET; ORAL

TRAMADOL HYDROCHLORIDE AND ACETAMINOPHEN

<u>AB</u>	MACLEODS PHARMS LTD	<u>325MG; 37.5MG</u>	<u>A206885 001</u>	May 02, 2017
<u>AB</u>	MICRO LABS LTD	<u>325MG; 37.5MG</u>	<u>A201952 001</u>	Dec 14, 2012
	INDIA			
<u>AB</u>	MYLAN	<u>325MG; 37.5MG</u>	<u>A077858 001</u>	Sep 26, 2008
<u>AB</u>	SUN PHARM INDS INC	<u>325MG; 37.5MG</u>	<u>A077184 001</u>	Dec 16, 2005
<u>AB</u>	ZYDUS PHARMS USA	<u>325MG; 37.5MG</u>	<u>A090460 001</u>	Sep 06, 2012
	INC			

ULTRACET

<u>AB</u>	+! JANSSEN PHARMS	<u>325MG; 37.5MG</u>	<u>N021123 001</u>	Aug 15, 2001
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ACETAZOLAMIDE

CAPSULE, EXTENDED RELEASE; ORAL

ACETAZOLAMIDE

<u>AB</u>	ALEMBIC PHARMS LTD	<u>500MG</u>	<u>A210423 001</u>	Feb 19, 2019
<u>AB</u>	CADILA	<u>500MG</u>	<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>	! HERITAGE PHARMS INC	<u>500MG</u>	<u>A090779 001</u>	Jul 14, 2011
<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>	EYWA PHARMA	<u>125MG</u>	<u>A211556 001</u>	Oct 18, 2019
<u>AB</u>		<u>250MG</u>	<u>A211556 002</u>	Oct 18, 2019
<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>	LANNETT	<u>250MG</u>	<u>A084840 001</u>	
<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>	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

ACETAZOLAMIDE SODIUM

INJECTABLE; INJECTION

ACETAZOLAMIDE SODIUM

<u>AP</u>	MYLAN ASI	<u>EQ 500MG BASE/VIAL</u>	<u>A200880 001</u>	May 09, 2012
<u>AP</u>	WEST-WARD PHARMS	<u>EQ 500MG BASE/VIAL</u>	<u>A040089 001</u>	Feb 28, 1995
	INT			
<u>AP</u>	! XGEN PHARMS	<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>	MLV	<u>2%</u>	<u>A040607 001</u>	Feb 24, 2005
<u>AT</u>	RISING	<u>2%</u>	<u>A207280 001</u>	Mar 09, 2018
<u>AT</u>	TARO	<u>2%</u>	<u>A088638 001</u>	Sep 06, 1984
<u>AT</u>	! WOCKHARDT BIO AG	<u>2%</u>	<u>A040166 001</u>	Jul 26, 1996

VOSOL

<u>AT</u>	HI TECH PHARMA	<u>2%</u>	<u>N012179 001</u>	
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ACETIC ACID, GLACIAL; HYDROCORTISONE

SOLUTION/DROPS; OTIC

HYDROCORTISONE AND ACETIC ACID

<u>AT</u>	TARO PHARM INDS LTD	<u>2%; 1%</u>	<u>A088759 001</u>	Mar 04, 1985
<u>AT</u>	VINTAGE	<u>2%; 1%</u>	<u>A040609 001</u>	Feb 06, 2006

VOSOL HC

<u>AT</u>	+! HI TECH PHARMA	<u>2%; 1%</u>	<u>N012770 001</u>	
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ACETOHYDROXAMIC ACID

TABLET; ORAL

LITHOSTAT

	+! MISSION PHARMA	<u>250MG</u>	<u>N018749 001</u>	May 31, 1983
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PRESCRIPTION DRUG PRODUCT LIST

ACETYLCHOLINE CHLORIDEFOR SOLUTION;OPHTHALMIC
MIOCHOL-E

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

ACETYLCYSTEINE

INJECTABLE; INTRAVENOUS

ACETADOTEAP +! CUMBERLAND PHARMS 6GM/30ML (200MG/ML) N021539 001 Jan 23, 2004ACETYLCYSTEINEAP AKORN INC 6GM/30ML (200MG/ML) A203173 001 Mar 24, 2015AP AUROBINDO PHARMA LTD 6GM/30ML (200MG/ML) A207358 001 Feb 29, 2016AP FRESENIUS KABI USA 6GM/30ML (200MG/ML) A200644 001 Nov 07, 2012AP SAGENT PHARMS INC 6GM/30ML (200MG/ML) A091684 001 Oct 31, 2017AP ZYDUS PHARMS 6GM/30ML (200MG/ML) A208166 001 Jul 20, 2018

SOLUTION;INHALATION, ORAL

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

CAPSULE;ORAL

ACITRETINAB BARR LABS INC 10MG A091455 001 Apr 04, 2013AB 25MG A091455 002 Apr 04, 2013AB IMPAX LABS INC 10MG A202552 001 Dec 23, 2015AB 17.5MG A202552 002 Dec 23, 2015AB 22.5MG A202552 003 Dec 23, 2015AB 25MG A202552 004 Dec 23, 2015AB MYLAN 10MG A202148 001 Sep 10, 2015AB 25MG A202148 002 Sep 10, 2015AB SIGMAPHARM LABS LLC 10MG A204633 001 May 22, 2015AB 17.5MG A204633 002 May 22, 2015AB 22.5MG A204633 003 May 22, 2015AB 25MG A204633 004 May 22, 2015AB TEVA PHARMS USA 17.5MG A202897 001 Apr 04, 2013AB 22.5MG A202897 002 Apr 04, 2013SORIATANEAB + STIEFEL LABS INC 10MG N019821 001 Oct 28, 1996AB +! 25MG N019821 002 Oct 28, 1996ACLIDINIUM BROMIDE

POWDER, METERED;INHALATION

TUDORZA PRESSAIR

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

ACLIDINIUM BROMIDE; FORMOTEROL FUMARATE

POWDER, METERED;INHALATION

DUAKLIR PRESSAIR

CIRCASSIA 0.4MG/INH;0.012MG/INH N210595 001 Mar 29, 2019

ACRIVASTINE; PSEUDOEPHEDRINE HYDROCHLORIDE

CAPSULE;ORAL

SEMPREX-D

+! AUXILIUM PHARMS LLC 8MG;60MG N019806 001 Mar 25, 1994

ACYCLOVIR

CAPSULE;ORAL

ACYCLOVIRAB ! APOTEX 200MG A075677 001 Sep 28, 2005AB CADILA 200MG A204313 001 Mar 25, 2016AB CADILA PHARMS LTD 200MG A201445 001 Mar 06, 2014AB CARLSBAD TECHNOLOGY 200MG A206261 001 Aug 16, 2017AB DAVA PHARMS INC 200MG A074833 001 Apr 22, 1997AB HERITAGE PHARMS INC 200MG A074889 001 Oct 31, 1997AB KENTON 200MG A075090 001 Jan 26, 1999AB TEVA 200MG A074578 001 Apr 22, 1997

CREAM;TOPICAL

ACYCLOVIRAB ! PERRIGO UK FINCO 5% A208702 001 Feb 04, 2019

PRESCRIPTION DRUG PRODUCT LIST

ACYCLOVIR

CREAM; TOPICAL

ZOVIRAX

AB	+	BAUSCH	5%	N021478 001	Dec 30, 2002
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OINTMENT; TOPICAL

ACYCLOVIR

AB		ACP NIMBLE	5%	A205591 001	Nov 13, 2017
AB		ALEMBIC PHARMS LTD	5%	A209000 001	Apr 06, 2018
AB		AMNEAL PHARMS	5%	A204605 001	Jun 18, 2014
AB		APOTEX	5%	A210774 001	Sep 06, 2019
AB		CADILA	5%	A205974 001	Mar 15, 2019
AB		CIPLA	5%	A211794 001	Jan 18, 2019
AB		FOUGERA PHARMS INC	5%	A206633 001	May 11, 2016
AB		GLENMARK PHARMS SA	5%	A205510 001	Jul 31, 2017
AB		MYLAN PHARMS INC	5%	A202459 001	Apr 03, 2013
AB		TARO	5%	A205469 001	Dec 21, 2016
AB		TOLMAR	5%	A206437 001	Jul 31, 2017
AB		TORRENT	5%	A209971 001	Jan 11, 2019
AB		XIROMED	5%	A201501 001	Jan 29, 2020

ZOVIRAX

AB	+	BAUSCH	5%	N018604 001	Mar 29, 1982
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SUSPENSION; ORAL

ACYCLOVIR

AB		ACTAVIS MID ATLANTIC	200MG/5ML	A074738 001	Apr 28, 1997
AB		HI TECH PHARMA	200MG/5ML	A077026 001	Jun 07, 2005

ZOVIRAX

AB	+	MYLAN	200MG/5ML	N019909 001	Dec 22, 1989
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TABLET; BUCCAL

SITAVIG

+	EPI HLTH	50MG	N203791 001	Apr 12, 2013
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TABLET; ORAL

ACYCLOVIR

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

ACYCLOVIR SODIUM

INJECTABLE; INJECTION

ACYCLOVIR SODIUM

AP		AUROBINDO PHARMA LTD	EQ 50MG BASE/ML	A203701 001	Oct 11, 2013
AP	!	FRESENIUS KABI USA	EQ 50MG BASE/ML	A074930 001	May 13, 1998
	!	ZYDUS PHARMS	EQ 500MG BASE/VIAL	A206535 001	Aug 31, 2018
	!		EQ 1GM BASE/VIAL	A206535 002	Aug 31, 2018

ACYCLOVIR; HYDROCORTISONE

CREAM; TOPICAL

XERESE

+	BAUSCH	5%; 1%	N022436 001	Jul 31, 2009
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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 **0.3%** **A201000 001** Oct 27, 2014

ATLANTIC

AB GLENMARK GENERICS **0.1%** **A091314 001** Jul 01, 2010

AB P AND L **0.1%** **A090962 001** Jun 02, 2010

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

AB TOLMAR **0.3%** **A200298 001** Jun 14, 2012

DIFFERIN

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

LOTION; TOPICAL

DIFFERIN

+! GALDERMA LABS LP 0.1%

N022502 001 Mar 17, 2010

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 MID **0.1%; 2.5%** **A203790 001** Sep 30, 2015

ATLANTIC

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

AB PERRIGO UK FINCO **0.1%; 2.5%** **A205033 001** Jan 23, 2018

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

AB TOLMAR **0.1%; 2.5%** **A206164 001** May 23, 2018

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

ADEFOVIR DIPIVOXIL

TABLET; ORAL

ADEFOVIR DIPIVOXIL

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

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

HEPSERA

AB +! GILEAD **10MG** **N021449 001** Sep 20, 2002

ADENOSINE

INJECTABLE; INJECTION

ADENOSINE

AP ! AKORN **3MG/ML** **A078076 001** Oct 31, 2008

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 MYLAN LABS LTD **3MG/ML** **A078640 001** Mar 21, 2014

AP **3MG/ML** **A078686 001** May 13, 2009

AP WEST-WARD PHARMS **3MG/ML** **A076404 001** Jun 16, 2004

INT

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

SOLUTION; INTRAVENOUS

ADENOSINE

AP AKORN **60MG/20ML (3MG/ML)** **A090450 001** Oct 02, 2014

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

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

LTD

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 HOSPIRA INC **60MG/20ML (3MG/ML)** **A203883 001** Mar 24, 2014

AP **90MG/30ML (3MG/ML)** **A203883 002** Mar 24, 2014

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

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

AP ! TEVA PHARMS USA **60MG/20ML (3MG/ML)** **A077425 001** Aug 29, 2013

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

PRESCRIPTION DRUG PRODUCT LIST

AFAMELANOTIDEIMPLANT; 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

ALBENDAZOLE**AB** ACTAVIS ELIZABETH **200MG** **A208094 001** May 20, 2019**AB** CIPLA LTD **200MG** **A210434 001** Sep 21, 2018**AB** EDENBRIDGE PHARMS **200MG** **A211117 001** May 14, 2019**AB** STRIDES PHARMA **200MG** **A210011 001** Dec 07, 2018**AB** ZYDUS PHARMS **200MG** **A208979 001** Dec 14, 2018ALBENZA**AB** +! IMPAX LABS INC **200MG** **N020666 001** Jun 11, 1996ALBUMIN HUMAN

INJECTABLE; INJECTION

OPTISON

+! GE HEALTHCARE 10MG/ML N020899 001 Dec 31, 1997

ALBUMIN IODINATED I-125 SERUM

INJECTABLE; INJECTION

JEANATOPE

+ ISO TEX 100uCi/10ML (10uCi/ML) N017836 003 Jun 08, 2004

+ 500uCi/0.5ML N017836 001

+! 1,000uCi/ML N017836 002

ALBUMIN IODINATED I-131 SERUM

INJECTABLE; INJECTION

MEGATOPE

+! ISO TEX 0.5mCi/VIAL N017837 001

+! 1mCi/VIAL N017837 002

ALBUTEROL SULFATE

AEROSOL, METERED; INHALATION

ALBUTEROL SULFATE**AB** PERRIGO PHARMS CO **EQ 0.09MG BASE/INH** **A203760 001** Feb 24, 2020PROAIR HFA**AB** +! TEVA BRANDED PHARM **EQ 0.09MG BASE/INH** **N021457 001** Oct 29, 2004

PROVENTIL-HFA

BX +! 3M DRUG DELIVERY EQ 0.09MG BASE/INH N020503 001 Aug 15, 1996

VENTOLIN HFA

BX +! GLAXOSMITHKLINE EQ 0.09MG BASE/INH N020983 001 Apr 19, 2001

POWDER, METERED; INHALATION

PROAIR DIGIHALER

+ TEVA BRANDED PHARM EQ 0.09MG BASE/INH N205636 002 Dec 21, 2018

PROAIR RESPICLICK

+! TEVA BRANDED PHARM EQ 0.09MG BASE/INH N205636 001 Mar 31, 2015

SOLUTION; INHALATION

ACCUNE**AN** +! MYLAN SPECIALITY LP **EQ 0.021% BASE** **N020949 002** Apr 30, 2001**AN** +! **EQ 0.042% BASE** **N020949 001** Apr 30, 2001ALBUTEROL SULFATE**AN** AUROBINDO PHARMA **EQ 0.083% BASE** **A206224 001** Oct 17, 2017

LTD

AN HI TECH PHARMA **EQ 0.5% BASE** **A074543 001** Jan 15, 1998**AN** NEPHRON **EQ 0.021% BASE** **A076355 002** Mar 31, 2010**AN** **EQ 0.042% BASE** **A076355 001** Jun 28, 2004**AN** ! **EQ 0.083% BASE** **A074880 001** Sep 17, 1997**AN** ! **EQ 0.5% BASE** **A075664 001** Jun 26, 2001**AN** RITEDOSE CORP **EQ 0.083% BASE** **A077839 001** Dec 16, 2008**AN** SUN PHARM **EQ 0.083% BASE** **A207857 001** Jul 21, 2017

PRESCRIPTION DRUG PRODUCT LIST

ALBUTEROL SULFATE

SOLUTION; INHALATION

ALBUTEROL SULFATE

AN	WATSON LABS	EQ 0.021% BASE	A077772 001	Sep 25, 2007
AN		EQ 0.042% BASE	A077772 002	Sep 25, 2007

SYRUP; ORAL

ALBUTEROL SULFATE

AA	AMNEAL PHARMS	EQ 2MG BASE/5ML	A079241 001	May 12, 2010
AA	COSETTE	EQ 2MG BASE/5ML	A074454 001	Sep 25, 1995
AA	HI TECH PHARMA	EQ 2MG BASE/5ML	A074749 001	Jan 30, 1998
AA	LANNETT CO INC	EQ 2MG BASE/5ML	A078105 001	Dec 27, 2006
AA	QUAGEN	EQ 2MG BASE/5ML	A212197 001	Sep 06, 2019
AA	! TEVA	EQ 2MG BASE/5ML	A073419 001	Mar 30, 1992
AA	VISTAPHARM	EQ 2MG BASE/5ML	A077788 001	Jun 26, 2007

TABLET; ORAL

ALBUTEROL SULFATE

AB	AMNEAL PHARMS CO	EQ 2MG BASE	A208804 001	May 21, 2018
AB		EQ 4MG BASE	A208804 002	May 21, 2018
AB	APPCO	EQ 2MG BASE	A207046 001	Jun 29, 2018
AB		EQ 4MG BASE	A207046 002	Jun 29, 2018
AB	ARISE	EQ 2MG BASE	A210948 001	Mar 15, 2019
AB		EQ 4MG BASE	A210948 002	Mar 15, 2019
AB	MYLAN	EQ 2MG BASE	A072894 002	Jan 17, 1991
AB	!	EQ 4MG BASE	A072894 001	Jan 17, 1991
AB	SUN PHARM INDUSTRIES	EQ 2MG BASE	A072637 002	Dec 05, 1989
AB		EQ 4MG BASE	A072637 001	Dec 05, 1989
AB	VIRTUS PHARM	EQ 2MG BASE	A211397 001	Oct 26, 2018
AB		EQ 4MG BASE	A211397 002	Oct 26, 2018

TABLET, EXTENDED RELEASE; ORAL

ALBUTEROL SULFATE

AB	MYLAN	EQ 4MG BASE	A078092 002	Jan 29, 2007
AB	VOSPIRE ER			
AB	DAVA PHARMS INC	EQ 4MG BASE	A076130 002	Sep 26, 2002
	ALBUTEROL SULFATE			
	! MYLAN	EQ 8MG BASE	A078092 001	Jan 29, 2007

ALBUTEROL SULFATE; IPRATROPIUM BROMIDE

SOLUTION; INHALATION

ALBUTEROL SULFATE AND IPRATROPIUM BROMIDE

AN	CIPLA	EQ 0.083% BASE; 0.017%	A077559 001	Dec 31, 2007
AN	NEPHRON	EQ 0.083% BASE; 0.017%	A076749 001	Dec 31, 2007
AN	RITEDOSE CORP	EQ 0.083% BASE; 0.017%	A202496 001	Oct 01, 2012
AN	SUN PHARM	EQ 0.083% BASE; 0.017%	A207875 001	Aug 07, 2017
AN	WATSON LABS TEVA	EQ 0.083% BASE; 0.017%	A077063 001	Dec 31, 2007

SPRAY, METERED; INHALATION

COMBIVENT RESPIMAT

+	! BOEHRINGER INGELHEIM	EQ 0.1MG BASE/INH; 0.02MG/INH	N021747 001	Oct 07, 2011
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ALCAFTADINE

SOLUTION/DROPS; OPHTHALMIC

LASTACFT

+	! ALLERGAN	0.25%	N022134 001	Jul 28, 2010
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ALCLOMETASONE DIPROPIONATE

CREAM; TOPICAL

ALCLOMETASONE DIPROPIONATE

AB	! FOUGERA PHARMS	0.05%	A076973 001	Jul 12, 2005
AB	GLENMARK GENERICS	0.05%	A079061 001	Jun 23, 2009
AB	TARO	0.05%	A076587 001	Sep 15, 2005

OINTMENT; TOPICAL

ALCLOMETASONE DIPROPIONATE

AB	! FOUGERA PHARMS	0.05%	A076884 001	Jul 18, 2005
AB	GLENMARK GENERICS	0.05%	A079227 001	Jul 30, 2009
AB	TARO	0.05%	A076730 001	Jul 29, 2004

ALCOHOL

SOLUTION; INTRA-ARTERIAL

ABLYSINOL

+	BELCHER	99% (1ML)	N207987 001	Jun 21, 2018
+	!	99% (5ML)	N207987 002	Jun 21, 2018

PRESCRIPTION DRUG PRODUCT LIST

ALECTINIB HYDROCHLORIDE

CAPSULE; ORAL

ALECENSA

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

ALENDRONATE SODIUM

SOLUTION; ORAL

ALENDRONATE SODIUM

! HIKMA EQ 70MG BASE/75ML A090520 001 Feb 25, 2013

TABLET; ORAL

ALENDRONATE SODIUM

AB	APOTEX	EQ 5MG BASE	A077982 001	Aug 04, 2008
AB		EQ 10MG BASE	A077982 002	Aug 04, 2008
AB		EQ 35MG BASE	A077982 003	Aug 04, 2008
AB		EQ 70MG BASE	A077982 004	Aug 04, 2008
AB	AUROBINDO PHARMA	EQ 10MG BASE	A090124 001	Aug 04, 2008
AB		EQ 35MG BASE	A090124 002	Aug 04, 2008
AB		EQ 70MG BASE	A090124 003	Aug 04, 2008
AB	CIPLA	EQ 5MG BASE	A076768 001	Aug 04, 2008
AB		EQ 10MG BASE	A076768 002	Aug 04, 2008
AB		EQ 35MG BASE	A076768 003	Aug 04, 2008
AB		EQ 40MG BASE	A076768 004	Aug 04, 2008
AB		EQ 70MG BASE	A076768 005	Aug 04, 2008
AB	HANGZHOU BINJIANG	EQ 5MG BASE	A090258 001	Sep 24, 2009
AB		EQ 10MG BASE	A090258 002	Sep 24, 2009
AB		EQ 35MG BASE	A090258 003	Sep 24, 2009
AB		EQ 70MG BASE	A090258 004	Sep 24, 2009
AB	IMPAX LABS INC	EQ 5MG BASE	A075710 001	Feb 06, 2008
AB		EQ 10MG BASE	A075710 002	Feb 06, 2008
AB		EQ 35MG BASE	A075710 003	Feb 06, 2008
AB		EQ 40MG BASE	A075710 004	Feb 06, 2008
AB		EQ 70MG BASE	A075710 005	Feb 06, 2008
AB	JUBILANT CADISTA	EQ 5MG BASE	A090557 001	Feb 18, 2010
AB		EQ 10MG BASE	A090557 002	Feb 18, 2010
AB		EQ 35MG BASE	A090557 003	Feb 18, 2010
AB		EQ 70MG BASE	A090557 004	Feb 18, 2010
AB	NEOPHARMA	EQ 5MG BASE	A079049 003	Aug 04, 2008
AB		EQ 10MG BASE	A079049 004	Aug 04, 2008
AB		EQ 35MG BASE	A079049 001	Aug 04, 2008
AB		EQ 70MG BASE	A079049 002	Aug 04, 2008
AB	SUN PHARM	EQ 5MG BASE	A090022 001	Sep 10, 2008
AB		EQ 10MG BASE	A090022 002	Sep 10, 2008
AB		EQ 35MG BASE	A090022 003	Sep 10, 2008
AB		EQ 70MG BASE	A090022 004	Sep 10, 2008
AB	WATSON LABS	EQ 35MG BASE	A076984 001	Aug 04, 2008
AB		EQ 40MG BASE	A076984 002	Aug 04, 2008
AB		EQ 70MG BASE	A076984 003	Aug 04, 2008

FOSAMAX

AB	+! MERCK AND CO INC	EQ 70MG BASE	N020560 005	Oct 20, 2000
TABLET, EFFERVESCENT; ORAL				
BINOSTO				
	+! ASCEND THERAPS US	EQ 70MG BASE	N202344 001	Mar 12, 2012

ALENDRONATE SODIUM; CHOLECALCIFEROL

TABLET; ORAL

FOSAMAX PLUS D

+ MERCK EQ 70MG BASE; 2,800 IU N021762 001 Apr 07, 2005

+! EQ 70MG BASE; 5,600 IU N021762 002 Apr 26, 2007

ALFENTANIL HYDROCHLORIDE

INJECTABLE; INJECTION

ALFENTA

AP	+! AKORN	EQ 0.5MG BASE/ML	N019353 001	Dec 29, 1986
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ALFENTANIL

AP	HOSPIRA	EQ 0.5MG BASE/ML	A075221 001	Oct 28, 1999
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ALFUZOSIN HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL

ALFUZOSIN HYDROCHLORIDE

AB	APOTEX INC	10MG	A079013 001	Jul 18, 2011
AB	AUROBINDO PHARMA LTD	10MG	A079060 001	Aug 30, 2012
AB	INVAGEN PHARMS	10MG	A090284 001	Jan 17, 2012
AB	SUN PHARM	10MG	A079057 001	Jul 18, 2011
AB	TORRENT PHARMS	10MG	A079054 001	Jul 18, 2011

PRESCRIPTION DRUG PRODUCT LIST

ALFUZOSIN HYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL

ALFUZOSIN HYDROCHLORIDE

<u>AB</u>	UNICHEM LABS LTD	<u>10MG</u>	<u>A203192</u>	<u>001</u>	Jan 28, 2016
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UROXATRAL

<u>AB</u>	+!	CONCORDIA	<u>10MG</u>	<u>N021287</u>	<u>001</u>	Jun 12, 2003
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ALISKIREN HEMIFUMARATE

TABLET;ORAL

ALISKIREN HEMIFUMARATE

<u>AB</u>	ANCHEN PHARMS	<u>EQ 150MG BASE</u>	<u>A206665</u>	<u>001</u>	Mar 22, 2019
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<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A206665</u>	<u>002</u>	Mar 22, 2019
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TEKTURNA

<u>AB</u>	+	NODEN PHARMA	<u>EQ 150MG BASE</u>	<u>N021985</u>	<u>001</u>	Mar 05, 2007
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<u>AB</u>	+		<u>EQ 300MG BASE</u>	<u>N021985</u>	<u>002</u>	Mar 05, 2007
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ALISKIREN HEMIFUMARATE; HYDROCHLOROTHIAZIDE

TABLET;ORAL

TEKTURNA HCT

+	NODEN PHARMA	EQ 150MG BASE;12.5MG	N022107	001	Jan 18, 2008
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+		EQ 150MG BASE;25MG	N022107	002	Jan 18, 2008
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+	!	EQ 300MG BASE;12.5MG	N022107	003	Jan 18, 2008
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+	!	EQ 300MG BASE;25MG	N022107	004	Jan 18, 2008
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ALITRETINOIN

GEL;TOPICAL

PANRETIN

+	!	CONCORDIA	EQ 0.1% BASE	N020886	001	Feb 02, 1999
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ALLOPURINOL

TABLET;ORAL

ALLOPURINOL

<u>AB</u>	ACCORD HLTHCARE	<u>100MG</u>	<u>A203154</u>	<u>001</u>	May 06, 2013
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<u>AB</u>		<u>300MG</u>	<u>A203154</u>	<u>002</u>	May 06, 2013
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<u>AB</u>	APOTEX INC	<u>100MG</u>	<u>A077353</u>	<u>001</u>	Sep 08, 2005
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<u>AB</u>		<u>300MG</u>	<u>A077353</u>	<u>002</u>	Sep 08, 2005
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<u>AB</u>	INDOCO REMEDIES	<u>100MG</u>	<u>A204467</u>	<u>001</u>	Jul 28, 2016
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<u>AB</u>		<u>300MG</u>	<u>A204467</u>	<u>002</u>	Jul 28, 2016
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<u>AB</u>	IPCA LABS LTD	<u>100MG</u>	<u>A090637</u>	<u>001</u>	Mar 16, 2011
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<u>AB</u>		<u>300MG</u>	<u>A090637</u>	<u>002</u>	Mar 16, 2011
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<u>AB</u>	MYLAN	<u>100MG</u>	<u>A018659</u>	<u>001</u>	Oct 24, 1986
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<u>AB</u>		<u>300MG</u>	<u>A018659</u>	<u>002</u>	Oct 24, 1986
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<u>AB</u>	NORTHSTAR HLTHCARE	<u>100MG</u>	<u>A078253</u>	<u>001</u>	Sep 11, 2007
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<u>AB</u>		<u>300MG</u>	<u>A078253</u>	<u>002</u>	Sep 11, 2007
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<u>AB</u>	SUN PHARM	<u>100MG</u>	<u>A071450</u>	<u>002</u>	Jan 09, 1987
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<u>AB</u>	INDUSTRIES				
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<u>AB</u>		<u>300MG</u>	<u>A071450</u>	<u>001</u>	Jan 09, 1987
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<u>AB</u>	UNICHEM LABS LTD	<u>100MG</u>	<u>A211820</u>	<u>001</u>	Mar 12, 2019
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<u>AB</u>		<u>300MG</u>	<u>A211820</u>	<u>002</u>	Mar 12, 2019
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<u>AB</u>	VINTAGE PHARMS	<u>100MG</u>	<u>A075798</u>	<u>001</u>	Jun 27, 2003
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<u>AB</u>		<u>300MG</u>	<u>A075798</u>	<u>002</u>	Jun 27, 2003
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<u>AB</u>	WATSON LABS	<u>100MG</u>	<u>N018832</u>	<u>002</u>	Sep 28, 1984
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<u>AB</u>		<u>300MG</u>	<u>N018877</u>	<u>001</u>	Sep 28, 1984
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<u>AB</u>	ZYDUS PHARMS	<u>100MG</u>	<u>A210117</u>	<u>001</u>	Oct 12, 2017
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<u>AB</u>		<u>300MG</u>	<u>A210117</u>	<u>002</u>	Oct 12, 2017
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LOPURIN

<u>AB</u>	DR REDDYS LA	<u>100MG</u>	<u>A071586</u>	<u>001</u>	Apr 02, 1987
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<u>AB</u>		<u>300MG</u>	<u>A071587</u>	<u>001</u>	Apr 02, 1987
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ZYLOPRIM

<u>AB</u>	+	CASPER PHARMA LLC	<u>100MG</u>	<u>N016084</u>	<u>001</u>
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<u>AB</u>	+	!	<u>300MG</u>	<u>N016084</u>	<u>002</u>
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ALLOPURINOL SODIUM

INJECTABLE; INJECTION

ALLOPURINOL SODIUM

<u>AP</u>	WEST-WARD PHARMS	<u>EQ 500MG BASE/VIAL</u>	<u>A076870</u>	<u>001</u>	Aug 26, 2004
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ALOPRIM

<u>AP</u>	+	MYLAN INSTITUTIONAL	<u>EQ 500MG BASE/VIAL</u>	<u>N020298</u>	<u>001</u>	May 17, 1996
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PRESCRIPTION DRUG PRODUCT LISTALMOTRIPTAN MALATE

TABLET; ORAL

ALMOTRIPTAN MALATE

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

OSENI

+	TAKEDA PHARMS USA	EQ 12.5MG BASE;EQ 15MG BASE	N022426 004	Jan 25, 2013
+		EQ 12.5MG BASE;EQ 30MG BASE	N022426 005	Jan 25, 2013
+		EQ 12.5MG BASE;EQ 45MG BASE	N022426 006	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

AB	AMNEAL PHARMS	EQ 0.5MG BASE	A206647 001	Dec 22, 2016
AB		EQ 1MG BASE	A206647 002	Dec 22, 2016
AB	EYWA PHARMA	EQ 0.5MG BASE	A211621 001	Sep 16, 2019
AB		EQ 1MG BASE	A211621 002	Sep 16, 2019
AB	HIKMA	EQ 0.5MG BASE	A200652 001	May 04, 2015
AB		EQ 1MG BASE	A200652 002	May 04, 2015
AB	PAR PHARM INC	EQ 0.5MG BASE	A206113 001	Feb 23, 2018
AB		EQ 1MG BASE	A206113 002	Feb 23, 2018
AB	RISING	EQ 0.5MG BASE	A209180 001	Jan 14, 2019
AB		EQ 1MG BASE	A209180 002	Jan 14, 2019

LOTRONEX

AB	+	SEBELA IRELAND LTD	EQ 0.5MG BASE	N021107 002	Dec 23, 2003
AB	+	!	EQ 1MG BASE	N021107 001	Feb 09, 2000

ALPELISIB

TABLET; ORAL

PIQRAY

+	NOVARTIS	50MG	N212526 001	May 24, 2019
+		150MG	N212526 002	May 24, 2019
+	!	200MG	N212526 003	May 24, 2019

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

PRESCRIPTION DRUG PRODUCT LIST

ALPRAZOLAM

CONCENTRATE; ORAL

ALPRAZOLAM

! HIKMA

1MG/ML

A074312 001 Oct 31, 1993

TABLET; ORAL

ALPRAZOLAM

<u>AB</u>	ACTAVIS ELIZABETH	<u>0.25MG</u>	<u>A074342 001</u>	Oct 31, 1993
<u>AB</u>		<u>0.5MG</u>	<u>A074342 002</u>	Oct 31, 1993
<u>AB</u>		<u>1MG</u>	<u>A074342 003</u>	Oct 31, 1993
<u>AB</u>		<u>2MG</u>	<u>A074342 004</u>	Oct 31, 1993
<u>AB</u>	APOTEX INC	<u>0.25MG</u>	<u>A077741 001</u>	Jan 19, 2007
<u>AB</u>		<u>0.5MG</u>	<u>A077741 002</u>	Jan 19, 2007
<u>AB</u>		<u>1MG</u>	<u>A077741 003</u>	Jan 19, 2007
<u>AB</u>		<u>2MG</u>	<u>A077741 004</u>	Jan 19, 2007
<u>AB</u>	AUROBINDO PHARMA LTD	<u>0.25MG</u>	<u>A203346 001</u>	Jul 31, 2015
<u>AB</u>		<u>0.5MG</u>	<u>A203346 002</u>	Jul 31, 2015
<u>AB</u>		<u>1MG</u>	<u>A203346 003</u>	Jul 31, 2015
<u>AB</u>		<u>2MG</u>	<u>A203346 004</u>	Jul 31, 2015
<u>AB</u>	BRECKENRIDGE	<u>0.25MG</u>	<u>A207507 001</u>	Jul 09, 2018
<u>AB</u>		<u>0.5MG</u>	<u>A207507 002</u>	Jul 09, 2018
<u>AB</u>		<u>1MG</u>	<u>A207507 003</u>	Jul 09, 2018
<u>AB</u>		<u>2MG</u>	<u>A207507 004</u>	Jul 09, 2018
<u>AB</u>	MYLAN	<u>0.25MG</u>	<u>A074215 001</u>	Jan 27, 1994
<u>AB</u>		<u>0.5MG</u>	<u>A074215 002</u>	Jan 27, 1994
<u>AB</u>		<u>1MG</u>	<u>A074215 003</u>	Jan 27, 1994
<u>AB</u>		<u>2MG</u>	<u>A074215 004</u>	Jan 27, 1994
<u>AB</u>	NATCO PHARMA LTD	<u>0.25MG</u>	<u>A200739 001</u>	Apr 15, 2015
<u>AB</u>		<u>0.5MG</u>	<u>A200739 002</u>	Apr 15, 2015
<u>AB</u>		<u>1MG</u>	<u>A200739 003</u>	Apr 15, 2015
<u>AB</u>		<u>2MG</u>	<u>A200739 004</u>	Apr 15, 2015
<u>AB</u>	OXFORD PHARMS	<u>0.25MG</u>	<u>A078491 001</u>	Sep 25, 2008
<u>AB</u>		<u>0.5MG</u>	<u>A078491 002</u>	Sep 25, 2008
<u>AB</u>		<u>1MG</u>	<u>A078491 003</u>	Sep 25, 2008
<u>AB</u>		<u>2MG</u>	<u>A078491 004</u>	Dec 12, 2008
<u>AB</u>	SANDOZ	<u>0.25MG</u>	<u>A074112 001</u>	Dec 29, 1995
<u>AB</u>		<u>0.5MG</u>	<u>A074112 002</u>	Dec 29, 1995
<u>AB</u>		<u>1MG</u>	<u>A074112 003</u>	Dec 29, 1995
<u>AB</u>		<u>2MG</u>	<u>A074909 001</u>	Mar 25, 1998
<u>AB</u>	SUN PHARM	<u>0.25MG</u>	<u>A090082 001</u>	Jun 17, 2010
<u>AB</u>		<u>0.5MG</u>	<u>A090082 002</u>	Jun 17, 2010
<u>AB</u>		<u>1MG</u>	<u>A090082 003</u>	Jun 17, 2010
<u>AB</u>		<u>2MG</u>	<u>A090082 004</u>	Jun 17, 2010
<u>AB</u>	VINTAGE PHARMS	<u>0.25MG</u>	<u>A090248 001</u>	Sep 17, 2010
<u>AB</u>		<u>0.5MG</u>	<u>A090248 002</u>	Sep 17, 2010
<u>AB</u>		<u>1MG</u>	<u>A090248 003</u>	Sep 17, 2010
<u>AB</u>		<u>2MG</u>	<u>A090248 004</u>	Sep 17, 2010

XANAX

<u>AB</u>	+ PHARMACIA AND 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>	Nov 27, 1985

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>	ANCHEN PHARMS	<u>0.5MG</u>	<u>A078469 001</u>	Sep 29, 2011
<u>AB</u>		<u>1MG</u>	<u>A078469 002</u>	Sep 29, 2011
<u>AB</u>		<u>2MG</u>	<u>A078469 003</u>	Sep 29, 2011
<u>AB</u>		<u>3MG</u>	<u>A078469 004</u>	Sep 29, 2011
<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 LTD	<u>0.5MG</u>	<u>A090871 001</u>	Jun 07, 2011

PRESCRIPTION DRUG PRODUCT LIST

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ALPRAZOLAM

TABLET, EXTENDED RELEASE;ORAL

ALPRAZOLAM

<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>	+	PHARMACIA AND 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>		PAR PHARM	<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>		TEVA PHARMS USA	<u>0.5MG/ML</u>	<u>A075196 001</u> Apr 30, 1999
<u>AP</u>		WEST-WARD PHARMS INT	<u>0.5MG/ML</u>	<u>A074815 001</u> Jan 20, 1998

CAVERJECT

<u>AP</u>	+	PHARMACIA AND UPJOHN	<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

EDEX

<u>AP</u>	+	AUXILIUM PHARMS LLC	<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

PROSTIN VR PEDIATRIC

<u>AP</u>	+	PHARMACIA AND UPJOHN	<u>0.5MG/ML</u>	<u>N018484 001</u>
CAVERJECT IMPULSE				
		PHARMACIA AND UPJOHN	0.01MG/VIAL	N021212 001 Jun 11, 2002
			0.02MG/VIAL	N021212 002 Jun 11, 2002
EDEX				
	+	AUXILIUM PHARMS LLC	0.01MG/VIAL	N020649 005 Jul 30, 1998
	+		0.02MG/VIAL	N020649 006 Jul 30, 1998
	+		0.04MG/VIAL	N020649 007 Jul 30, 1998
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

ALVIMOPAN

CAPSULE; ORAL

ALVIMOPAN

<u>AB</u>		WATSON LABS TEVA	<u>12MG</u>	<u>A208295 001</u> Dec 19, 2019
<u>ENTEREG</u>				
<u>AB</u>	+	CUBIST PHARMS	<u>12MG</u>	<u>N021775 001</u> May 20, 2008

AMANTADINE HYDROCHLORIDE

CAPSULE; ORAL

AMANTADINE HYDROCHLORIDE

<u>AB</u>		ALEMBIC PHARMS LTD	<u>100MG</u>	<u>A208966 001</u> Jun 21, 2017
<u>AB</u>		BIONPHARMA INC	<u>100MG</u>	<u>A078720 001</u> May 29, 2008
<u>AB</u>		HERITAGE PHARMA	<u>100MG</u>	<u>A209171 001</u> Jun 12, 2017
<u>AB</u>		INVAGEN PHARMS	<u>100MG</u>	<u>A207570 001</u> Sep 30, 2016
<u>AB</u>		INVATECH	<u>100MG</u>	<u>A210129 001</u> Mar 02, 2020
<u>AB</u>	!	SANDOZ	<u>100MG</u>	<u>A071293 001</u> Feb 18, 1987
<u>AB</u>		STRIDES PHARMA	<u>100MG</u>	<u>A209047 001</u> Jun 07, 2017
<u>AB</u>		UPSHER SMITH LABS	<u>100MG</u>	<u>A070589 001</u> Aug 05, 1986
<u>AB</u>		WATSON LABS INC	<u>100MG</u>	<u>A208107 001</u> Dec 06, 2016

PRESCRIPTION DRUG PRODUCT LIST

AMANTADINE HYDROCHLORIDE

CAPSULE;ORAL

AMANTADINE HYDROCHLORIDE

AB	!	ZYDUS PHARMS	100MG	A208278	001	May 31, 2016
CAPSULE, EXTENDED RELEASE;ORAL						
GOCOVRI						
	+	ADAMAS PHARMA	EQ 68.5MG BASE	N208944	001	Aug 24, 2017
	+	!	EQ 137MG BASE	N208944	002	Aug 24, 2017

SYRUP;ORAL

AMANTADINE HYDROCHLORIDE

AA	!	ANDA REPOSITORY	50MG/5ML	A074028	001	Jun 28, 1993
AA	!	CMP PHARMA INC	50MG/5ML	A075819	001	Sep 11, 2002
AA	!	HI TECH PHARMA	50MG/5ML	A074170	001	Oct 28, 1994
AA	!	PHARM ASSOC	50MG/5ML	A074509	001	Jul 17, 1995
AA	!	WOCKHARDT BIO AG	50MG/5ML	A075060	001	Dec 24, 1998

TABLET;ORAL

AMANTADINE HYDROCHLORIDE

AB		INVAGEN PHARMS	100MG	A207571	001	Jan 31, 2017
AB		INVATECH	100MG	A210215	001	Mar 10, 2020
AB		JUBILANT GENERICS	100MG	A210403	001	Feb 07, 2018
AB		STRIDES PHARMA	100MG	A209035	001	Jun 09, 2017
AB	!	UPSHER SMITH LABS	100MG	A076186	001	Dec 16, 2002
AB		WATSON LABS INC	100MG	A208096	001	Dec 15, 2016
TABLET, EXTENDED RELEASE;ORAL						
OSMOLEX ER						
	+	OSMOTICA PHARM	EQ 129MG BASE	N209410	001	Feb 16, 2018
	+		EQ 193MG BASE	N209410	002	Feb 16, 2018
	+	!	EQ 258MG BASE	N209410	003	Feb 16, 2018

AMBRISENTAN

TABLET;ORAL

AMBRISENTAN

AB		CIPLA	5MG	A210715	001	Apr 26, 2019
AB			10MG	A210715	002	Apr 26, 2019
AB		MYLAN	5MG	A208441	001	Mar 28, 2019
AB			10MG	A208441	002	Mar 28, 2019
AB		PAR PHARM INC	5MG	A209509	001	Apr 10, 2019
AB			10MG	A209509	002	Apr 10, 2019
AB		SIGMAPHARM LABS LLC	5MG	A208354	001	Apr 10, 2019
AB			10MG	A208354	002	Apr 10, 2019
AB		SUN PHARM	5MG	A210784	001	Mar 28, 2019
AB			10MG	A210784	002	Mar 28, 2019
AB		WATSON LABS INC	5MG	A208252	001	Mar 28, 2019
AB			10MG	A208252	002	Mar 28, 2019
AB		ZYDUS PHARMS	5MG	A210058	001	Mar 28, 2019
AB			10MG	A210058	002	Mar 28, 2019

LETAIRIS

AB	+	GILEAD	5MG	N022081	001	Jun 15, 2007
AB	+	!	10MG	N022081	002	Jun 15, 2007

AMCINONIDE

CREAM;TOPICAL

AMCINONIDE

AB	!	FOUGERA PHARMS	0.1%	A076065	001	May 15, 2003
AB		TARO PHARM INDS	0.1%	A076229	001	May 31, 2002

LOTION;TOPICAL

AMCINONIDE

	!	FOUGERA PHARMS	0.1%	A076329	001	Nov 06, 2002
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OINTMENT;TOPICAL

AMCINONIDE

AB	!	FOUGERA PHARMS	0.1%	A076096	001	Nov 19, 2002
AB		TARO PHARM INDS	0.1%	A076367	001	Mar 19, 2003

AMIFAMPRIDINE

TABLET;ORAL

RUZURGI

	+	JACOBUS PHARM CO INC	10MG	N209321	001	May 06, 2019
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PRESCRIPTION DRUG PRODUCT LIST

AMIFAMPRIDINE PHOSPHATE

TABLET; ORAL

FIRDAPSE

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

AMIFOSTINE

INJECTABLE; INJECTION

AMIFOSTINEAP ! MYLAN LABS LTD 500MG/VIAL A204363 001 Jul 17, 2017AP SUN PHARM 500MG/VIAL A077126 001 Mar 14, 2008ETHYOLAP +! CLINIGEN 500MG/VIAL N020221 001 Dec 08, 1995AMIKACIN SULFATE

INJECTABLE; INJECTION

AMIKACIN SULFATEAP ! EMCURE PHARMS LTD EQ 250MG BASE/ML A204040 001 Dec 12, 2013AP FRESENIUS KABI USA EQ 50MG BASE/ML A205605 001 Dec 09, 2015AP EQ 250MG BASE/ML A205604 001 Dec 09, 2015AP SAGENT PHARMS INC EQ 250MG BASE/ML A203323 001 May 12, 2016AP ! WEST-WARD PHARMS EQ 50MG BASE/ML A063313 001 Apr 11, 1994

INT

AP EQ 250MG BASE/ML A063315 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 HYDROCHLORIDEAB ! PAR PHARM 5MG A070346 001 Jan 22, 1986AB SIGMAPHARM LABS LLC 5MG A079133 001 Jan 30, 2009AB WINDLAS HLTHCARE 5MG A204180 001 Aug 07, 2015MIDAMORAB + PADDOCK LLC 5MG N018200 001AMILORIDE HYDROCHLORIDE; HYDROCHLOROTHIAZIDE

TABLET; ORAL

AMILORIDE HYDROCHLORIDE AND HYDROCHLOROTHIAZIDEAB BARR EQ 5MG ANHYDROUS; 50MG A071111 001 May 10, 1988AB ! MYLAN 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

FREAMINE HBC 6.9%

B BRAUN 6.9% (6.9GM/100ML) N016822 006 May 17, 1983

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

NEPHRAMINE 5.4%

B BRAUN 5.4% (5.4GM/100ML) N017766 001

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

PRESCRIPTION DRUG PRODUCT LISTAMINO ACIDS

INJECTABLE; INJECTION

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 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 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
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;261MG/100ML;217MG/100ML;112MG/100ML	N020678 001	Mar 26, 1997
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;261MG/100ML;297MG/100ML;77MG/100ML	N020678 009	Mar 26, 1997
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;261MG/100ML;297MG/100ML;77MG/100ML	N020678 011	Mar 26, 1997
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;261MG/100ML;297MG/100ML;77MG/100ML	N020678 012	Mar 26, 1997
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;261MG/100ML;297MG/100ML;77MG/100ML	N020678 008	Mar 26, 1997
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;261MG/100ML;340MG/100ML;59MG/100ML	N020678 016	Mar 26, 1997
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;261MG/100ML;340MG/100ML;59MG/100ML	N020678 017	Mar 26, 1997
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;261MG/100ML;340MG/100ML;59MG/100ML	N020678 018	Mar 26, 1997
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;261MG/100ML;340MG/100ML;59MG/100ML	N020678 019	Mar 26, 1997
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;261MG/100ML;340MG/100ML;59MG/100ML	N020678 021	Mar 26, 1997

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

PRESCRIPTION DRUG PRODUCT LIST

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

EMULSION; INTRAVENOUS

PERIKABIVEN IN PLASTIC CONTAINER

+	2.4%; 20MG/100ML; 6.8GM/100ML; 68MG/100ML; 124MG/100ML; 170MG/100ML; 105MG/100ML; 3.5GM/100ML (1920ML)	N200656 002	Aug 25, 2014
+!	2.4%; 20MG/100ML; 6.8GM/100ML; 68MG/100ML; 124MG/100ML; 170MG/100ML; 105MG/100ML; 3.5GM/100ML (2400ML)	N200656 003	Aug 25, 2014

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
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

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; 440MG/100ML; 690MG/100ML	N016822 007	Jul 01, 1988
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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; 200MG/100ML; 120MG/100ML	N016822 003	
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AMINOCAPROIC ACID

INJECTABLE; INJECTION

<u>AP</u>	<u>AMINOCAPROIC ACID</u>	<u>250MG/ML</u>	<u>A071192 001</u>	Dec 01, 1987
	LUITPOLD			
<u>AP</u>	<u>AMINOCAPROIC ACID IN PLASTIC CONTAINER</u>	<u>250MG/ML</u>	<u>A070010 001</u>	Mar 09, 1987
	HOSPIRA			
	SYRUP; ORAL			
<u>AA</u>	<u>AMICAR</u>	<u>1.25GM/5ML</u>	<u>N015230 002</u>	
	CLOVER PHARMS			
<u>AA</u>	<u>AMINOCAPROIC ACID</u>	<u>1.25GM/5ML</u>	<u>A212780 001</u>	Aug 23, 2019
	AMNEAL PHARMS LLC			
<u>AA</u>	<u>AMINOCAPROIC ACID</u>	<u>1.25GM/5ML</u>	<u>A212814 001</u>	Feb 26, 2020
	VISTAPHARM			
	TABLET; ORAL			
<u>AB</u>	<u>AMICAR</u>	<u>500MG</u>	<u>N015197 001</u>	
	CLOVER PHARMS			
<u>AB</u>	<u>AMICAR</u>	<u>1GM</u>	<u>N015197 002</u>	Jun 24, 2004
	CLOVER PHARMS			
<u>AB</u>	<u>AMINOCAPROIC ACID</u>	<u>500MG</u>	<u>A212492 001</u>	Nov 26, 2019
	AMNEAL			
<u>AB</u>	<u>AMINOCAPROIC ACID</u>	<u>500MG</u>	<u>A209060 001</u>	Nov 27, 2018
	SUNNY PHARMTECH INC			
<u>AB</u>	<u>AMINOCAPROIC ACID</u>	<u>1GM</u>	<u>A209060 002</u>	Nov 27, 2018
	SUNNY PHARMTECH INC			

PRESCRIPTION DRUG PRODUCT LISTAMINOLEVULINIC 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					
+!	DUSA	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					
!	JACOBUS	4GM/PACKET	A074346	001	Jun 30, 1994

AMIODARONE HYDROCHLORIDE

INJECTABLE;INJECTION

AMIODARONE HYDROCHLORIDE

AP	AUROBINDO PHARMA LTD	50MG/ML	A204550	001	Oct 25, 2017	
AP	!	FRESENIUS KABI USA	50MG/ML	A075761	001	Oct 15, 2002
AP	!	GLAND PHARMA LTD	50MG/ML	A077161	001	Apr 20, 2005
AP		HIKMA FARMACEUTICA	50MG/ML	A077234	001	Feb 25, 2008
AP		HOSPIRA INC	50MG/ML	A203884	001	Nov 25, 2013
AP			50MG/ML	A203885	001	Nov 25, 2013
AP	!	MYLAN INSTITUTIONAL	50MG/ML	A076217	001	Oct 15, 2002
AP		WOCKHARDT	50MG/ML	A077610	001	Oct 30, 2008
AP			50MG/ML	A077834	001	Oct 30, 2008
	NEXTERONE					
+!	BAKTER 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

AB	AUROBINDO PHARMA LTD	200MG	A204742	001	Jun 03, 2016	
AB		100MG	A075389	002	Dec 28, 2017	
AB	MAYNE PHARMA INC	200MG	A075389	001	Jan 25, 2001	
AB		400MG	A075389	003	Dec 28, 2017	
AB	MURTY PHARMS	100MG	A077069	003	Oct 04, 2016	
AB		200MG	A077069	001	Apr 08, 2005	
AB		400MG	A077069	002	Apr 08, 2005	
AB	RUBICON	200MG	A078578	001	Nov 06, 2008	
AB	!	SANDOZ	200MG	A075315	001	Dec 23, 1998
AB		400MG	A075315	002	Jun 30, 2000	
AB	TARO	100MG	A075424	002	Dec 18, 2002	
AB		200MG	A075424	001	Mar 30, 2001	
AB		400MG	A076362	001	Nov 29, 2002	
AB	TEVA PHARMS	200MG	A074739	001	Nov 30, 1998	
AB	ZYDUS PHARMS USA INC	200MG	A079029	001	Sep 16, 2008	
	PACERONE					
AB	UPSHER SMITH LABS	100MG	A075135	002	Apr 12, 2005	
AB		200MG	A075135	001	Apr 30, 1998	
	AMIODARONE HYDROCHLORIDE					
	TARO	300MG	A076362	002	Dec 02, 2003	

AMISULPRIDE

SOLUTION;INTRAVENOUS

BARHEMSYS					
+!	ACACIA PHARMA LTD	5MG/2ML (2.5MG/ML)	N209510	001	Feb 26, 2020

AMITRIPTYLINE HYDROCHLORIDE

TABLET;ORAL

AMITRIPTYLINE HYDROCHLORIDE

AB	ACCORD HLTHCARE	10MG	A202446	001	Jun 04, 2014
AB		25MG	A202446	002	Jun 04, 2014
AB		50MG	A202446	003	Jun 04, 2014
AB		75MG	A202446	004	Jun 04, 2014
AB		100MG	A202446	005	Jun 04, 2014

PRESCRIPTION DRUG PRODUCT LIST

AMITRIPTYLINE HYDROCHLORIDE

TABLET;ORAL

AMITRIPTYLINE HYDROCHLORIDE

<u>AB</u>		<u>150MG</u>	<u>A202446 006</u>	Jun 04, 2014
<u>AB</u>	MYLAN	<u>10MG</u>	<u>A086009 002</u>	
<u>AB</u>		<u>25MG</u>	<u>A086009 003</u>	
<u>AB</u>		<u>50MG</u>	<u>A086009 001</u>	
<u>AB</u>		<u>75MG</u>	<u>A086009 004</u>	
<u>AB</u>		<u>100MG</u>	<u>A086009 005</u>	
<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>75MG</u>	<u>A085971 001</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>	VINTAGE PHARMS	<u>10MG</u>	<u>A040218 001</u>	Sep 11, 1997
<u>AB</u>		<u>25MG</u>	<u>A040218 002</u>	Sep 11, 1997
<u>AB</u>		<u>50MG</u>	<u>A040218 003</u>	Sep 11, 1997
<u>AB</u>		<u>75MG</u>	<u>A040218 004</u>	Sep 11, 1997
<u>AB</u>		<u>100MG</u>	<u>A040218 005</u>	Sep 11, 1997
<u>AB</u>		<u>150MG</u>	<u>A040218 006</u>	Sep 11, 1997
<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

	MYLAN PHARMS INC	EQ 12.5MG BASE;5MG	A071297 002	Dec 10, 1986
	!	EQ 25MG BASE;10MG	A071297 001	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

	+!	SILVERGATE PHARMS	EQ 1MG BASE/ML	N211340 001	Jul 08, 2019
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AMLODIPINE BESYLATE

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
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A078925 003</u>	May 04, 2009
<u>AB</u>	APOTEX	<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>	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>	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

PRESCRIPTION DRUG PRODUCT LIST

AMLODIPINE BESYLATE

TABLET;ORAL

AMLODIPINE BESYLATE

<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>	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>	HEBEI CHANGSHAN	<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>	HIKMA PHARMS	<u>EQ 2.5MG BASE</u>	<u>A077771 001</u>	Apr 12, 2011
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A077771 002</u>	Apr 12, 2011
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A077771 003</u>	Apr 12, 2011
<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>	MACLEODS PHARMS LTD	<u>EQ 5MG BASE</u>	<u>A201380 001</u>	Apr 13, 2012
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A201380 002</u>	Apr 13, 2012
<u>AB</u>	MYLAN	<u>EQ 2.5MG BASE</u>	<u>A076418 001</u>	Oct 03, 2005
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A076418 002</u>	Oct 03, 2005
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A076418 003</u>	Oct 03, 2005
<u>AB</u>	ORCHID HLTHCARE	<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>	SUN PHARM INDS LTD	<u>EQ 2.5MG BASE</u>	<u>A077974 001</u>	Jul 09, 2007
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A077974 002</u>	Jul 09, 2007
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A077974 003</u>	Jul 09, 2007
<u>AB</u>	TORRENT PHARMS	<u>EQ 2.5MG BASE</u>	<u>A078573 001</u>	Sep 22, 2008
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A078573 002</u>	Sep 22, 2008
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A078573 003</u>	Sep 22, 2008
<u>AB</u>	UNICHEM LABS LTD	<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>	UPSHER SMITH LABS	<u>EQ 2.5MG BASE</u>	<u>A077759 001</u>	Jul 09, 2007
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A077759 002</u>	Jul 09, 2007
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A077759 003</u>	Jul 09, 2007
<u>AB</u>	WATSON LABS	<u>EQ 2.5MG BASE</u>	<u>A077671 001</u>	Jul 19, 2007
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A077671 002</u>	Jul 19, 2007
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A077671 003</u>	Jul 19, 2007
<u>AB</u>	WOCKHARDT	<u>EQ 2.5MG BASE</u>	<u>A078500 001</u>	Sep 06, 2007
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A078500 002</u>	Sep 06, 2007
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A078500 003</u>	Sep 06, 2007
<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>	+ PFIZER	<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

PRESCRIPTION DRUG PRODUCT LIST

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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	
<u>AB</u>		<u>EQ 5MG BASE;EQ 40MG BASE</u>	<u>A203874 006</u>	Mar 07, 2014	
<u>AB</u>		<u>EQ 5MG BASE;EQ 80MG BASE</u>	<u>A203874 007</u>	Mar 07, 2014	
<u>AB</u>		<u>EQ 10MG BASE;EQ 10MG BASE</u>	<u>A203874 008</u>	Mar 07, 2014	
<u>AB</u>		<u>EQ 10MG BASE;EQ 20MG BASE</u>	<u>A203874 009</u>	Mar 07, 2014	
<u>AB</u>		<u>EQ 10MG BASE;EQ 40MG BASE</u>	<u>A203874 010</u>	Mar 07, 2014	
<u>AB</u>		<u>EQ 10MG BASE;EQ 80MG BASE</u>	<u>A203874 011</u>	Mar 07, 2014	
<u>AB</u>	MYLAN	<u>EQ 2.5MG BASE;EQ 10MG BASE</u>	<u>A200465 001</u>	Nov 29, 2013	
<u>AB</u>		<u>EQ 2.5MG BASE;EQ 20MG BASE</u>	<u>A200465 002</u>	Nov 29, 2013	
<u>AB</u>		<u>EQ 2.5MG BASE;EQ 40MG BASE</u>	<u>A200465 003</u>	Nov 29, 2013	
<u>AB</u>		<u>EQ 5MG BASE;EQ 10MG BASE</u>	<u>A200465 004</u>	Nov 29, 2013	
<u>AB</u>		<u>EQ 5MG BASE;EQ 20MG BASE</u>	<u>A200465 005</u>	Nov 29, 2013	
<u>AB</u>		<u>EQ 5MG BASE;EQ 40MG BASE</u>	<u>A200465 006</u>	Nov 29, 2013	
<u>AB</u>		<u>EQ 5MG BASE;EQ 80MG BASE</u>	<u>A200465 007</u>	Nov 29, 2013	
<u>AB</u>		<u>EQ 10MG BASE;EQ 10MG BASE</u>	<u>A200465 008</u>	Nov 29, 2013	
<u>AB</u>		<u>EQ 10MG BASE;EQ 20MG BASE</u>	<u>A200465 009</u>	Nov 29, 2013	
<u>AB</u>		<u>EQ 10MG BASE;EQ 40MG BASE</u>	<u>A200465 010</u>	Nov 29, 2013	
<u>AB</u>		<u>EQ 10MG BASE;EQ 80MG BASE</u>	<u>A200465 011</u>	Nov 29, 2013	
<u>AB</u>	ZYDUS PHARMS	<u>EQ 2.5MG BASE;EQ 10MG BASE</u>	<u>A207762 001</u>	Jan 11, 2019	
<u>AB</u>		<u>EQ 2.5MG BASE;EQ 20MG BASE</u>	<u>A207762 002</u>	Jan 11, 2019	
<u>AB</u>		<u>EQ 2.5MG BASE;EQ 40MG BASE</u>	<u>A207762 003</u>	Jan 11, 2019	
<u>AB</u>		<u>EQ 5MG BASE;EQ 10MG BASE</u>	<u>A207762 004</u>	Jan 11, 2019	
<u>AB</u>		<u>EQ 5MG BASE;EQ 20MG BASE</u>	<u>A207762 005</u>	Jan 11, 2019	
<u>AB</u>		<u>EQ 5MG BASE;EQ 40MG BASE</u>	<u>A207762 006</u>	Jan 11, 2019	
<u>AB</u>		<u>EQ 5MG BASE;EQ 80MG BASE</u>	<u>A207762 007</u>	Jan 11, 2019	
<u>AB</u>		<u>EQ 10MG BASE;EQ 10MG BASE</u>	<u>A207762 008</u>	Jan 11, 2019	
<u>AB</u>		<u>EQ 10MG BASE;EQ 20MG BASE</u>	<u>A207762 009</u>	Jan 11, 2019	
<u>AB</u>		<u>EQ 10MG BASE;EQ 40MG BASE</u>	<u>A207762 010</u>	Jan 11, 2019	
<u>AB</u>		<u>EQ 10MG BASE;EQ 80MG BASE</u>	<u>A207762 011</u>	Jan 11, 2019	
<u>CADUET</u>					
<u>AB</u>	+	PFIZER	<u>EQ 2.5MG BASE;EQ 10MG BASE</u>	<u>N021540 009</u>	Jul 29, 2004
<u>AB</u>	+		<u>EQ 2.5MG BASE;EQ 20MG BASE</u>	<u>N021540 010</u>	Jul 29, 2004
<u>AB</u>	+		<u>EQ 2.5MG BASE;EQ 40MG BASE</u>	<u>N021540 011</u>	Jul 29, 2004
<u>AB</u>	+		<u>EQ 5MG BASE;EQ 10MG BASE</u>	<u>N021540 001</u>	Jan 30, 2004
<u>AB</u>	+		<u>EQ 5MG BASE;EQ 20MG BASE</u>	<u>N021540 002</u>	Jan 30, 2004
<u>AB</u>	+		<u>EQ 5MG BASE;EQ 40MG BASE</u>	<u>N021540 003</u>	Jan 30, 2004
<u>AB</u>	+		<u>EQ 5MG BASE;EQ 80MG BASE</u>	<u>N021540 004</u>	Jan 30, 2004
<u>AB</u>	+		<u>EQ 10MG BASE;EQ 10MG BASE</u>	<u>N021540 005</u>	Jan 30, 2004
<u>AB</u>	+		<u>EQ 10MG BASE;EQ 20MG BASE</u>	<u>N021540 006</u>	Jan 30, 2004
<u>AB</u>	+		<u>EQ 10MG BASE;EQ 40MG BASE</u>	<u>N021540 007</u>	Jan 30, 2004
<u>AB</u>	+		<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

<u>AB</u>	APOTEX	<u>EQ 2.5MG BASE;10MG</u>	<u>A091431 001</u>	Dec 30, 2013
<u>AB</u>		<u>EQ 5MG BASE;10MG</u>	<u>A091431 002</u>	Dec 30, 2013
<u>AB</u>		<u>EQ 5MG BASE;20MG</u>	<u>A091431 003</u>	Dec 30, 2013
<u>AB</u>		<u>EQ 5MG BASE;40MG</u>	<u>A091431 004</u>	Dec 30, 2013
<u>AB</u>		<u>EQ 10MG BASE;20MG</u>	<u>A091431 005</u>	Dec 30, 2013
<u>AB</u>		<u>EQ 10MG BASE;40MG</u>	<u>A091431 006</u>	Dec 30, 2013
<u>AB</u>	AUROBINDO PHARMA LTD	<u>EQ 2.5MG BASE;10MG</u>	<u>A202239 001</u>	Sep 05, 2012
<u>AB</u>		<u>EQ 5MG BASE;10MG</u>	<u>A202239 002</u>	Sep 05, 2012
<u>AB</u>		<u>EQ 5MG BASE;20MG</u>	<u>A202239 003</u>	Sep 05, 2012
<u>AB</u>		<u>EQ 5MG BASE;40MG</u>	<u>A202239 004</u>	Sep 05, 2012
<u>AB</u>		<u>EQ 10MG BASE;20MG</u>	<u>A202239 005</u>	Sep 05, 2012
<u>AB</u>		<u>EQ 10MG BASE;40MG</u>	<u>A202239 006</u>	Sep 05, 2012

PRESCRIPTION DRUG PRODUCT LIST

AMLODIPINE BESYLATE; BENAZEPRIL HYDROCHLORIDE

CAPSULE;ORAL

AMLODIPINE BESYLATE AND BENAZEPRIL HYDROCHLORIDE

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	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	PAR PHARM	<u>EQ 2.5MG BASE;10MG</u>	<u>A078381 001</u>	Jul 29, 2010
AB		<u>EQ 5MG BASE;10MG</u>	<u>A078381 002</u>	Jul 29, 2010
AB		<u>EQ 5MG BASE;20MG</u>	<u>A078381 003</u>	Jul 29, 2010
AB		<u>EQ 5MG BASE;40MG</u>	<u>A078381 005</u>	Jul 29, 2010
AB		<u>EQ 10MG BASE;20MG</u>	<u>A078381 004</u>	Jul 29, 2010
AB		<u>EQ 10MG BASE;40MG</u>	<u>A078381 006</u>	Jul 29, 2010
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
AB	WATSON LABS INC	<u>EQ 5MG BASE;40MG</u>	<u>A090364 001</u>	Jul 05, 2011
AB		<u>EQ 10MG BASE;40MG</u>	<u>A090364 002</u>	Jul 05, 2011
<u>LOTREL</u>				
AB	+	NOVARTIS	<u>EQ 2.5MG BASE;10MG</u>	<u>N020364 002</u> Mar 03, 1995
AB	+		<u>EQ 5MG BASE;10MG</u>	<u>N020364 003</u> Mar 03, 1995
AB	+		<u>EQ 5MG BASE;20MG</u>	<u>N020364 004</u> Mar 03, 1995
AB	+		<u>EQ 5MG BASE;40MG</u>	<u>N020364 007</u> Apr 11, 2006
AB	+		<u>EQ 10MG BASE;20MG</u>	<u>N020364 005</u> Jun 20, 2002
AB	+	!	<u>EQ 10MG BASE;40MG</u>	<u>N020364 006</u> Apr 11, 2006

AMLODIPINE BESYLATE; CELECOXIB

TABLET;ORAL

CONSENSI

+	COEPTIS	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; OLMESARTAN MEDOXOMIL

TABLET;ORAL

OLMESARTAN MEDOXOMIL, AMLODIPINE AND HYDROCHLOROTHIAZIDE

AB	PAR PHARM INC	<u>EQ 5MG BASE;12.5MG;20MG</u>	<u>A206137 001</u>	Oct 26, 2016
AB		<u>EQ 5MG BASE;12.5MG;40MG</u>	<u>A206137 002</u>	Oct 26, 2016
AB		<u>EQ 5MG BASE;25MG;40MG</u>	<u>A206137 003</u>	Oct 26, 2016
AB		<u>EQ 10MG BASE;12.5MG;40MG</u>	<u>A206137 004</u>	Oct 26, 2016
AB		<u>EQ 10MG BASE;25MG;40MG</u>	<u>A206137 005</u>	Oct 26, 2016
AB	TORRENT	<u>EQ 5MG BASE;12.5MG;20MG</u>	<u>A203580 001</u>	Oct 26, 2016
AB		<u>EQ 5MG BASE;12.5MG;40MG</u>	<u>A203580 002</u>	Oct 26, 2016
AB		<u>EQ 5MG BASE;25MG;40MG</u>	<u>A203580 003</u>	Oct 26, 2016
AB		<u>EQ 10MG BASE;12.5MG;40MG</u>	<u>A203580 004</u>	Oct 26, 2016
AB		<u>EQ 10MG BASE;25MG;40MG</u>	<u>A203580 005</u>	Oct 26, 2016
<u>TRIBENZOR</u>				
AB	+	DAIICHI SANKYO	<u>EQ 5MG BASE;12.5MG;20MG</u>	<u>N200175 001</u> Jul 23, 2010
AB	+		<u>EQ 5MG BASE;12.5MG;40MG</u>	<u>N200175 002</u> Jul 23, 2010
AB	+		<u>EQ 5MG BASE;25MG;40MG</u>	<u>N200175 003</u> Jul 23, 2010
AB	+		<u>EQ 10MG BASE;12.5MG;40MG</u>	<u>N200175 004</u> Jul 23, 2010
AB	+	!	<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

AB	AUROBINDO PHARMA LTD	<u>5MG;12.5MG;160MG</u>	<u>A206180 001</u>	Dec 19, 2017
AB		<u>5MG;25MG;160MG</u>	<u>A206180 002</u>	Dec 19, 2017
AB		<u>10MG;12.5MG;160MG</u>	<u>A206180 003</u>	Dec 19, 2017
AB		<u>10MG;25MG;160MG</u>	<u>A206180 004</u>	Dec 19, 2017
AB		<u>10MG;25MG;320MG</u>	<u>A206180 005</u>	Dec 19, 2017
AB	LUPIN LTD	<u>5MG;12.5MG;160MG</u>	<u>A200797 001</u>	Jun 03, 2015
AB		<u>5MG;25MG;160MG</u>	<u>A200797 002</u>	Jun 03, 2015
AB		<u>10MG;12.5MG;160MG</u>	<u>A200797 003</u>	Jun 03, 2015

PRESCRIPTION DRUG PRODUCT LIST

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AMLODIPINE BESYLATE; HYDROCHLOROTHIAZIDE; VALSARTAN

TABLET; ORAL

AMLODIPINE BESYLATE, VALSARTAN AND HYDROCHLOROTHIAZIDE

<u>AB</u>		<u>10MG;25MG;160MG</u>	<u>A200797 004</u>	Jun 03, 2015
<u>AB</u>		<u>10MG;25MG;320MG</u>	<u>A200797 005</u>	Jun 03, 2015
<u>AB</u>	PAR PHARM	<u>5MG;12.5MG;160MG</u>	<u>A201087 001</u>	Jun 01, 2015
<u>AB</u>		<u>5MG;25MG;160MG</u>	<u>A201087 002</u>	Jun 01, 2015
<u>AB</u>		<u>10MG;12.5MG;160MG</u>	<u>A201087 003</u>	Jun 01, 2015
<u>AB</u>		<u>10MG;25MG;160MG</u>	<u>A201087 004</u>	Jun 01, 2015
<u>AB</u>		<u>10MG;25MG;320MG</u>	<u>A201087 005</u>	Jun 01, 2015
<u>EXFORGE HCT</u>				
<u>AB</u>	+	NOVARTIS	<u>5MG;12.5MG;160MG</u>	<u>N022314 001</u> Apr 30, 2009
<u>AB</u>	+		<u>5MG;25MG;160MG</u>	<u>N022314 002</u> Apr 30, 2009
<u>AB</u>	+		<u>10MG;12.5MG;160MG</u>	<u>N022314 003</u> Apr 30, 2009
<u>AB</u>	+		<u>10MG;25MG;160MG</u>	<u>N022314 004</u> Apr 30, 2009
<u>AB</u>	+		<u>10MG;25MG;320MG</u>	<u>N022314 005</u> Apr 30, 2009

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>	ALEMIC PHARMS LTD	<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 LTD	<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>	GLENMARK PHARMS LTD	<u>EQ 5MG BASE;20MG</u>	<u>A207807 001</u>	Jul 05, 2017
<u>AB</u>		<u>EQ 5MG BASE;40MG</u>	<u>A207807 002</u>	Jul 05, 2017
<u>AB</u>		<u>EQ 10MG BASE;20MG</u>	<u>A207807 003</u>	Jul 05, 2017
<u>AB</u>		<u>EQ 10MG BASE;40MG</u>	<u>A207807 004</u>	Jul 05, 2017
<u>AB</u>	JUBILANT GENERICS	<u>EQ 5MG BASE;20MG</u>	<u>A207450 001</u>	May 15, 2017
<u>AB</u>		<u>EQ 5MG BASE;40MG</u>	<u>A207450 002</u>	May 15, 2017
<u>AB</u>		<u>EQ 10MG BASE;20MG</u>	<u>A207450 003</u>	May 15, 2017
<u>AB</u>		<u>EQ 10MG BASE;40MG</u>	<u>A207450 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>	TORRENT	<u>EQ 5MG BASE;20MG</u>	<u>A202933 001</u>	Nov 25, 2016
<u>AB</u>		<u>EQ 5MG BASE;40MG</u>	<u>A202933 002</u>	Nov 25, 2016
<u>AB</u>		<u>EQ 10MG BASE;20MG</u>	<u>A202933 003</u>	Nov 25, 2016
<u>AB</u>		<u>EQ 10MG BASE;40MG</u>	<u>A202933 004</u>	Nov 25, 2016
<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>	+	DAIICHI SANKYO	<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

PRESCRIPTION DRUG PRODUCT LIST

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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

AMLODIPINE BESYLATE; TELMISARTAN

TABLET; ORAL

TELMISARTAN AND AMLODIPINE

<u>AB</u>	ALEMBIC PHARMS LTD	<u>EQ 5MG BASE;40MG</u>	<u>A205234 001</u>	Nov 17, 2016
<u>AB</u>		<u>EQ 5MG BASE;80MG</u>	<u>A205234 003</u>	Nov 17, 2016
<u>AB</u>		<u>EQ 10MG BASE;40MG</u>	<u>A205234 002</u>	Nov 17, 2016
<u>AB</u>		<u>EQ 10MG BASE;80MG</u>	<u>A205234 004</u>	Nov 17, 2016
<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
<u>AB</u>	TORRENT	<u>EQ 5MG BASE;40MG</u>	<u>A202517 001</u>	Jan 08, 2014
<u>AB</u>		<u>EQ 5MG BASE;80MG</u>	<u>A202517 003</u>	Jan 08, 2014
<u>AB</u>		<u>EQ 10MG BASE;40MG</u>	<u>A202517 002</u>	Jan 08, 2014
<u>AB</u>		<u>EQ 10MG BASE;80MG</u>	<u>A202517 004</u>	Jan 08, 2014

TWYNSTA

<u>AB</u>	+	BOEHRINGER INGELHEIM	<u>EQ 5MG BASE;40MG</u>	<u>N022401 001</u>	Oct 16, 2009
<u>AB</u>	+		<u>EQ 5MG BASE;80MG</u>	<u>N022401 003</u>	Oct 16, 2009
<u>AB</u>	+		<u>EQ 10MG BASE;40MG</u>	<u>N022401 002</u>	Oct 16, 2009
<u>AB</u>	+	!	<u>EQ 10MG BASE;80MG</u>	<u>N022401 004</u>	Oct 16, 2009

AMLODIPINE BESYLATE; VALSARTAN

TABLET; ORAL

AMLODIPINE BESYLATE AND VALSARTAN

<u>AB</u>	ALEMBIC PHARMS LTD	<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 LTD	<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>	INVAGEN PHARMS	<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>	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>	PAR PHARM INC	<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

EXFORGE

<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

PRESCRIPTION DRUG PRODUCT LIST

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 HEALTH 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>	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>	PETNET	<u>30mCi-300mCi (3.75-37.5mCi/ML)</u>	<u>A204510 001</u>	Nov 02, 2015
<u>AP</u>	SOFIE	<u>18.8mCi-188mCi/5ML (3.75-37.5mCi/ML)</u>	<u>A204667 001</u>	Apr 22, 2015
<u>AP</u>	SPECTRON MRC LLC	<u>30mCi-300mCi/8ML (3.75-37.5mCi/ML)</u>	<u>A204455 001</u>	Apr 23, 2015
<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 TX SW MEDCTR	<u>30mCi-300mCi/8ML (3.75-37.5mCi/ML)</u>	<u>A209507 001</u>	Nov 01, 2019
<u>AP</u>	WA UNIV SCH MED	<u>30mCi-300mCi/8ML (3.75-37.5mCi/ML)</u>	<u>A204506 001</u>	Feb 07, 2014
	ESSENTIAL ISOTOPES	3.75-260mCi/ML	A205687 001	Dec 17, 2015
	NCM USA BRONX LLC	3.75-260mCi/mL	A204515 001	Feb 04, 2015
	PRECISION NUCLEAR	3.75-260mCi/ML	A204547 001	Aug 14, 2015
	SHERTECH LABS LLC	3.75-260mCi/ML	A204366 001	Sep 19, 2014
	SOFIE	3.75-260mCi/ML	A203543 001	Dec 14, 2012
	WI MEDCL CYCLOTRON	3.75-260mCi/ML	A204356 001	Dec 18, 2014

AMMONIUM CHLORIDE

INJECTABLE; INJECTION

AMMONIUM CHLORIDE IN PLASTIC CONTAINER

!	HOSPIRA	5MEQ/ML	A088366 001	Jun 13, 1984
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AMMONIUM LACTATE

CREAM; TOPICAL

AMMONIUM LACTATE

<u>AB</u>	! PERRIGO 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
<u>AB</u>	WATSON LABS INC	<u>EQ 12% BASE</u>	<u>A076829 001</u>	Feb 07, 2006

LOTION; TOPICAL

AMMONIUM LACTATE

<u>AB</u>	! PERRIGO 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
<u>AB</u>	WATSON LABS INC	<u>EQ 12% BASE</u>	<u>A075575 001</u>	Jun 11, 2002

AMOXAPINE

TABLET; ORAL

AMOXAPINE

	WATSON LABS	25MG	A072691 002	Aug 28, 1992
		50MG	A072691 003	Aug 28, 1992
		100MG	A072691 004	Aug 28, 1992
!		150MG	A072691 001	Aug 28, 1992

AMOXICILLIN

CAPSULE; ORAL

AMOXICILLIN

<u>AB</u>	AM ANTIBIOTICS	<u>250MG</u>	<u>A062058 001</u>	
<u>AB</u>		<u>500MG</u>	<u>A062058 002</u>	
<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>	DAVA PHARMS INC	<u>250MG</u>	<u>A062884 001</u>	Feb 25, 1988
<u>AB</u>		<u>500MG</u>	<u>A062881 001</u>	Feb 25, 1988
<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>	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>	NEOPHARMA	<u>250MG</u>	<u>A062216 001</u>	
<u>AB</u>		<u>500MG</u>	<u>A062216 004</u>	

PRESCRIPTION DRUG PRODUCT LIST

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AMOXICILLIN

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>	DAVA PHARMS INC	<u>125MG/5ML</u>	<u>A062927 001</u>	Nov 25, 1988
<u>AB</u>		<u>250MG/5ML</u>	<u>A062927 002</u>	Nov 25, 1988
<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
<u>AB</u>	WOCKHARDT BIO AG	<u>400MG/5ML</u>	<u>A065319 002</u>	Jun 18, 2007

AMOXICILLIN PEDIATRIC

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

<u>AB</u>	NEOPHARMA	<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>	

LAROTID

<u>AB</u>	NEOPHARMA	<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

AMOXICILLIN; CLARITHROMYCIN; LANSOPRAZOLE

CAPSULE, CAPSULE, DELAYED REL PELLETS, TABLET;ORAL

LANSOPRAZOLE, AMOXICILLIN AND CLARITHROMYCIN

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

AMOXICILLIN; CLARITHROMYCIN; OMEPRAZOLE

CAPSULE, TABLET, CAPSULE, DELAYED RELEASE;ORAL

OMEPRAZOLE AND CLARITHROMYCIN AND AMOXICILLIN

	+	CUMBERLAND PHARMS	500MG,N/A,N/A;N/A,500MG,N/A;N/A,N/A,20MG	N050824 001	Feb 08, 2011
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AMOXICILLIN; CLAVULANATE POTASSIUM

FOR SUSPENSION;ORAL

AMOXICILLIN AND CLAVULANATE POTASSIUM

<u>AB</u>	AUROBINDO PHARMA LTD	<u>125MG/5ML;EQ 31.25MG BASE/5ML</u>	<u>A209371 001</u>	Apr 19, 2019
<u>AB</u>		<u>200MG/5ML;EQ 28.5MG BASE/5ML</u>	<u>A201090 001</u>	Dec 20, 2011
<u>AB</u>		<u>250MG/5ML;EQ 62.5MG BASE/5ML</u>	<u>A209371 002</u>	Apr 19, 2019
<u>AB</u>		<u>400MG/5ML;EQ 57MG BASE/5ML</u>	<u>A201090 002</u>	Dec 20, 2011
<u>AB</u>		<u>600MG/5ML;EQ 42.9MG BASE/5ML</u>	<u>A201091 001</u>	Dec 20, 2011
<u>AB</u>	HIKMA PHARMS	<u>200MG/5ML;EQ 28.5MG BASE/5ML</u>	<u>A065191 002</u>	Jan 25, 2005
<u>AB</u>		<u>400MG/5ML;EQ 57MG BASE/5ML</u>	<u>A065191 001</u>	Jan 25, 2005
<u>AB</u>		<u>600MG/5ML;EQ 42.9MG BASE/5ML</u>	<u>A065373 001</u>	Nov 09, 2007
<u>AB</u>	SANDOZ	<u>200MG/5ML;EQ 28.5MG BASE/5ML</u>	<u>A065066 001</u>	Jun 05, 2002
<u>AB</u>		<u>400MG/5ML;EQ 57MG BASE/5ML</u>	<u>A065066 002</u>	Jun 05, 2002
<u>AB</u>	SANDOZ INC	<u>200MG/5ML;EQ 28.5MG BASE/5ML</u>	<u>A065098 001</u>	Dec 16, 2002
<u>AB</u>		<u>400MG/5ML;EQ 57MG BASE/5ML</u>	<u>A065098 002</u>	Dec 16, 2002

PRESCRIPTION DRUG PRODUCT LIST

AMOXICILLIN; CLAVULANATE POTASSIUM

FOR SUSPENSION; ORAL

AMOXICILLIN AND CLAVULANATE POTASSIUM

<u>AB</u>		<u>600MG/5ML;EQ 42.9MG BASE/5ML</u>	<u>A065358 001</u>	Aug 13, 2007
<u>AB</u>	TEVA	<u>200MG/5ML;EQ 28.5MG BASE/5ML</u>	<u>A065089 001</u>	May 25, 2004
<u>AB</u>	!	<u>400MG/5ML;EQ 57MG BASE/5ML</u>	<u>A065089 002</u>	May 25, 2004
<u>AB</u>	!	<u>600MG/5ML;EQ 42.9MG BASE/5ML</u>	<u>A065162 001</u>	Mar 12, 2004
<u>AB</u>	WOCKHARDT BIO AG	<u>250MG/5ML;EQ 62.5MG BASE/5ML</u>	<u>A065431 001</u>	Nov 25, 2008
<u>AB</u>		<u>600MG/5ML;EQ 42.9MG BASE/5ML</u>	<u>A065420 001</u>	Dec 02, 2013

AUGMENTIN '125'

<u>AB</u>	+	NEOPHARMA	<u>125MG/5ML;EQ 31.25MG BASE/5ML</u>	<u>N050575 001</u>	Aug 06, 1984
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AUGMENTIN '250'

<u>AB</u>	+	NEOPHARMA	<u>250MG/5ML;EQ 62.5MG BASE/5ML</u>	<u>N050575 002</u>	Aug 06, 1984
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TABLET; ORAL

AMOXICILLIN AND CLAVULANATE POTASSIUM

<u>AB</u>	AUROBINDO PHARMA LTD	<u>250MG;EQ 125MG BASE</u>	<u>A091569 001</u>	Jan 20, 2012	
<u>AB</u>		<u>500MG;EQ 125MG BASE</u>	<u>A091569 002</u>	Jan 20, 2012	
<u>AB</u>		<u>875MG;EQ 125MG BASE</u>	<u>A091568 001</u>	Jan 20, 2012	
<u>AB</u>	HIKMA PHARMS	<u>875MG;EQ 125MG BASE</u>	<u>A203824 001</u>	Aug 23, 2016	
<u>AB</u>	MICRO LABS LTD INDIA	<u>250MG;EQ 125MG BASE</u>	<u>A205707 001</u>	Dec 30, 2016	
<u>AB</u>		<u>500MG;EQ 125MG BASE</u>	<u>A205707 002</u>	Dec 30, 2016	
<u>AB</u>		<u>875MG;EQ 125MG BASE</u>	<u>A204755 003</u>	Dec 30, 2016	
<u>AB</u>	!	SANDOZ	<u>250MG;EQ 125MG BASE</u>	<u>A065189 001</u>	Aug 23, 2005
<u>AB</u>			<u>500MG;EQ 125MG BASE</u>	<u>A065064 001</u>	Mar 15, 2002
<u>AB</u>	!		<u>875MG;EQ 125MG BASE</u>	<u>A065063 001</u>	Mar 14, 2002
<u>AB</u>	!	SANDOZ INC	<u>500MG;EQ 125MG BASE</u>	<u>A065117 001</u>	Nov 27, 2002
<u>AB</u>			<u>875MG;EQ 125MG BASE</u>	<u>A065093 001</u>	Nov 21, 2002
<u>AB</u>	TEVA	<u>500MG;EQ 125MG BASE</u>	<u>A065101 001</u>	Oct 30, 2002	
<u>AB</u>	TEVA PHARMS USA	<u>875MG;EQ 125MG BASE</u>	<u>A065096 001</u>	Oct 29, 2002	

AUGMENTIN '875'

<u>AB</u>	+	NEOPHARMA	<u>875MG;EQ 125MG BASE</u>	<u>N050720 001</u>	Feb 13, 1996
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TABLET, CHEWABLE; ORAL

AMOXICILLIN AND CLAVULANATE POTASSIUM

	TEVA	<u>200MG;EQ 28.5MG BASE</u>	<u>A065205 001</u>	Feb 09, 2005
	!	<u>400MG;EQ 57MG BASE</u>	<u>A065205 002</u>	Feb 09, 2005

TABLET, EXTENDED RELEASE; ORAL

AMOXICILLIN AND CLAVULANATE POTASSIUM

<u>AB</u>	SANDOZ	<u>1GM;EQ 62.5MG BASE</u>	<u>A090227 001</u>	Apr 21, 2010
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AUGMENTIN XR

<u>AB</u>	+	NEOPHARMA	<u>1GM;EQ 62.5MG BASE</u>	<u>N050785 001</u>	Sep 25, 2002
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AMOXICILLIN; OMEPRAZOLE MAGNESIUM; RIFABUTIN

CAPSULE, DELAYED RELEASE; ORAL

TALICIA

+	!	REDHILL	<u>250MG;EQ 10MG BASE;12.5MG</u>	<u>N213004 001</u>	Nov 01, 2019
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AMPHETAMINE

SUSPENSION, EXTENDED RELEASE; ORAL

ADZENYS ER

+	!	NEOS THERAPS INC	<u>EQ 1.25MG BASE/ML</u>	<u>N204325 001</u>	Sep 15, 2017
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DYANAVEL XR

+	!	TRIS PHARMA INC	<u>EQ 2.5MG BASE/ML</u>	<u>N208147 001</u>	Oct 19, 2015
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TABLET, ORALLY DISINTEGRATING, EXTENDED RELEASE; ORAL

ADZENYS XR-ODT

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

AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATE

CAPSULE, EXTENDED RELEASE; ORAL

ADDERALL XR 10

<u>AB</u>	+	SHIRE	<u>2.5MG;2.5MG;2.5MG;2.5MG</u>	<u>N021303 001</u>	Oct 11, 2001
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ADDERALL XR 15

<u>AB</u>	+	SHIRE	<u>3.75MG;3.75MG;3.75MG;3.75MG</u>	<u>N021303 006</u>	May 22, 2002
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ADDERALL XR 20

<u>AB</u>	+	SHIRE	<u>5MG;5MG;5MG;5MG</u>	<u>N021303 002</u>	Oct 11, 2001
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ADDERALL XR 25

<u>AB</u>	+	SHIRE	<u>6.25MG;6.25MG;6.25MG;6.25MG</u>	<u>N021303 004</u>	May 22, 2002
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PRESCRIPTION DRUG PRODUCT LIST

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AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATE

CAPSULE, EXTENDED RELEASE; ORAL

ADDERALL XR 30

AB	+	SHIRE	7.5MG;7.5MG;7.5MG;7.5MG	N021303	003	Oct 11, 2001
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ADDERALL XR 5

AB	+	SHIRE	1.25MG;1.25MG;1.25MG;1.25MG	N021303	005	May 22, 2002
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DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE

AB		ACTAVIS ELIZABETH	1.25MG;1.25MG;1.25MG;1.25MG	A077302	001	Jun 22, 2012
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AB			2.5MG;2.5MG;2.5MG;2.5MG	A077302	002	Jun 22, 2012
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AB			3.75MG;3.75MG;3.75MG;3.75MG	A077302	003	Jun 22, 2012
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AB			5MG;5MG;5MG;5MG	A077302	004	Jun 22, 2012
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AB			6.25MG;6.25MG;6.25MG;6.25MG	A077302	005	Jun 22, 2012
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AB			7.5MG;7.5MG;7.5MG;7.5MG	A077302	006	Jun 22, 2012
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AB		ANI PHARMS INC	1.25MG;1.25MG;1.25MG;1.25MG	A205401	001	Jan 22, 2019
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AB			2.5MG;2.5MG;2.5MG;2.5MG	A205401	002	Jan 22, 2019
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AB			3.75MG;3.75MG;3.75MG;3.75MG	A205401	003	Jan 22, 2019
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AB			5MG;5MG;5MG;5MG	A205401	004	Jan 22, 2019
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AB			6.25MG;6.25MG;6.25MG;6.25MG	A205401	005	Jan 22, 2019
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AB			7.5MG;7.5MG;7.5MG;7.5MG	A205401	006	Jan 22, 2019
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AB		IMPAX LABS	1.25MG;1.25MG;1.25MG;1.25MG	A076852	001	Feb 16, 2016
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AB			2.5MG;2.5MG;2.5MG;2.5MG	A076852	002	Feb 16, 2016
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AB			3.75MG;3.75MG;3.75MG;3.75MG	A076852	003	Feb 16, 2016
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AB			5MG;5MG;5MG;5MG	A076852	004	Feb 16, 2016
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AB			6.25MG;6.25MG;6.25MG;6.25MG	A076852	005	Feb 16, 2016
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AB			7.5MG;7.5MG;7.5MG;7.5MG	A076852	006	Feb 16, 2016
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AB		PAR PHARM INC	1.25MG;1.25MG;1.25MG;1.25MG	A206159	001	May 31, 2019
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AB			2.5MG;2.5MG;2.5MG;2.5MG	A206159	002	May 31, 2019
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AB			3.75MG;3.75MG;3.75MG;3.75MG	A206159	003	May 31, 2019
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AB			5MG;5MG;5MG;5MG	A206159	004	May 31, 2019
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AB			6.25MG;6.25MG;6.25MG;6.25MG	A206159	005	May 31, 2019
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AB			7.5MG;7.5MG;7.5MG;7.5MG	A206159	006	May 31, 2019
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AB		RHODES PHARMS	1.25MG;1.25MG;1.25MG;1.25MG	A210651	001	May 17, 2019
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AB			2.5MG;2.5MG;2.5MG;2.5MG	A210651	002	May 17, 2019
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AB			3.75MG;3.75MG;3.75MG;3.75MG	A210651	003	May 17, 2019
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AB			5MG;5MG;5MG;5MG	A210651	004	May 17, 2019
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AB			6.25MG;6.25MG;6.25MG;6.25MG	A210651	005	May 17, 2019
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AB			7.5MG;7.5MG;7.5MG;7.5MG	A210651	006	May 17, 2019
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AB		SPECGX LLC	1.25MG;1.25MG;1.25MG;1.25MG	A211547	001	Apr 22, 2019
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AB			2.5MG;2.5MG;2.5MG;2.5MG	A211547	002	Apr 22, 2019
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AB			3.75MG;3.75MG;3.75MG;3.75MG	A211547	003	Apr 22, 2019
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AB			5MG;5MG;5MG;5MG	A211547	004	Apr 22, 2019
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AB			6.25MG;6.25MG;6.25MG;6.25MG	A211547	005	Apr 22, 2019
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AB			7.5MG;7.5MG;7.5MG;7.5MG	A211547	006	Apr 22, 2019
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AB		SUNGEN PHARMA	1.25MG;1.25MG;1.25MG;1.25MG	A212037	001	Dec 11, 2019
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AB			2.5MG;2.5MG;2.5MG;2.5MG	A212037	002	Dec 11, 2019
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AB			3.75MG;3.75MG;3.75MG;3.75MG	A212037	003	Dec 11, 2019
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AB			5MG;5MG;5MG;5MG	A212037	004	Dec 11, 2019
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AB			6.25MG;6.25MG;6.25MG;6.25MG	A212037	005	Dec 11, 2019
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AB			7.5MG;7.5MG;7.5MG;7.5MG	A212037	006	Dec 11, 2019
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AB		TEVA	1.25MG;1.25MG;1.25MG;1.25MG	A077488	001	Apr 29, 2013
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AB			2.5MG;2.5MG;2.5MG;2.5MG	A077488	002	Apr 29, 2013
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AB			3.75MG;3.75MG;3.75MG;3.75MG	A077488	003	Apr 29, 2013
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AB			5MG;5MG;5MG;5MG	A077488	004	Apr 29, 2013
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AB			6.25MG;6.25MG;6.25MG;6.25MG	A077488	005	Apr 29, 2013
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AB			7.5MG;7.5MG;7.5MG;7.5MG	A077488	006	Apr 29, 2013
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DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE

AB		BARR LABS INC	1.25MG;1.25MG;1.25MG;1.25MG	A076536	001	Feb 12, 2013
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AB			2.5MG;2.5MG;2.5MG;2.5MG	A076536	002	Feb 12, 2013
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AB			3.75MG;3.75MG;3.75MG;3.75MG	A076536	003	Feb 12, 2013
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AB			5MG;5MG;5MG;5MG	A076536	004	Feb 12, 2013
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AB			6.25MG;6.25MG;6.25MG;6.25MG	A076536	005	Feb 12, 2013
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AB			7.5MG;7.5MG;7.5MG;7.5MG	A076536	006	Feb 12, 2013
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MYDAYIS

	+	SHIRE DEV LLC	3.125MG;3.125MG;3.125MG;3.125MG	N022063	001	Jun 20, 2017
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	+		6.25MG;6.25MG;6.25MG;6.25MG	N022063	002	Jun 20, 2017
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	+		9.375MG;9.375MG;9.375MG;9.375MG	N022063	003	Jun 20, 2017
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	+		12.5MG;12.5MG;12.5MG;12.5MG	N022063	004	Jun 20, 2017
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TABLET; ORAL

DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE

AB		ACTAVIS ELIZABETH	1.25MG;1.25MG;1.25MG;1.25MG	A206340	001	Feb 05, 2016
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AB			1.875MG;1.875MG;1.875MG;1.875MG	A206340	002	Feb 05, 2016
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AB			2.5MG;2.5MG;2.5MG;2.5MG	A206340	003	Feb 05, 2016
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PRESCRIPTION DRUG PRODUCT LIST

AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATE

TABLET; ORAL

DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE

AB		<u>3.125MG;3.125MG;3.125MG;3.125MG</u>	<u>A206340 004</u>	Feb 05, 2016
AB		<u>3.75MG;3.75MG;3.75MG;3.75MG</u>	<u>A206340 005</u>	Feb 05, 2016
AB		<u>5MG;5MG;5MG;5MG</u>	<u>A206340 006</u>	Feb 05, 2016
AB		<u>7.5MG;7.5MG;7.5MG;7.5MG</u>	<u>A206340 007</u>	Feb 05, 2016
AB	ALVOGEN	<u>1.25MG;1.25MG;1.25MG;1.25MG</u>	<u>A207388 001</u>	Jul 28, 2017
AB		<u>1.875MG;1.875MG;1.875MG;1.875MG</u>	<u>A207388 002</u>	Jul 28, 2017
AB		<u>2.5MG;2.5MG;2.5MG;2.5MG</u>	<u>A207388 003</u>	Jul 28, 2017
AB		<u>3.125MG;3.125MG;3.125MG;3.125MG</u>	<u>A207388 004</u>	Jul 28, 2017
AB		<u>3.75MG;3.75MG;3.75MG;3.75MG</u>	<u>A207388 005</u>	Jul 28, 2017
AB		<u>5MG;5MG;5MG;5MG</u>	<u>A207388 006</u>	Jul 28, 2017
AB		<u>7.5MG;7.5MG;7.5MG;7.5MG</u>	<u>A207388 007</u>	Jul 28, 2017
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	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	MYLAN	<u>1.25MG;1.25MG;1.25MG;1.25MG</u>	<u>A206721 001</u>	Nov 10, 2015
AB		<u>1.875MG;1.875MG;1.875MG;1.875MG</u>	<u>A206721 002</u>	Nov 10, 2015
AB		<u>2.5MG;2.5MG;2.5MG;2.5MG</u>	<u>A206721 003</u>	Nov 10, 2015
AB		<u>3.125MG;3.125MG;3.125MG;3.125MG</u>	<u>A206721 004</u>	Nov 10, 2015
AB		<u>3.75MG;3.75MG;3.75MG;3.75MG</u>	<u>A206721 005</u>	Nov 10, 2015
AB		<u>5MG;5MG;5MG;5MG</u>	<u>A206721 006</u>	Nov 10, 2015
AB		<u>7.5MG;7.5MG;7.5MG;7.5MG</u>	<u>A206721 007</u>	Nov 10, 2015
AB	NESHER PHARMS	<u>1.25MG;1.25MG;1.25MG;1.25MG</u>	<u>A207340 001</u>	Oct 31, 2017
AB		<u>1.875MG;1.875MG;1.875MG;1.875MG</u>	<u>A207340 002</u>	Oct 31, 2017
AB		<u>2.5MG;2.5MG;2.5MG;2.5MG</u>	<u>A207340 003</u>	Oct 31, 2017
AB		<u>3.125MG;3.125MG;3.125MG;3.125MG</u>	<u>A207340 004</u>	Oct 31, 2017
AB		<u>3.75MG;3.75MG;3.75MG;3.75MG</u>	<u>A207340 005</u>	Oct 31, 2017
AB		<u>5MG;5MG;5MG;5MG</u>	<u>A207340 006</u>	Oct 31, 2017
AB		<u>7.5MG;7.5MG;7.5MG;7.5MG</u>	<u>A207340 007</u>	Oct 31, 2017
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	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
AB		<u>3.125MG;3.125MG;3.125MG;3.125MG</u>	<u>A040440 004</u>	Oct 07, 2003
AB		<u>3.75MG;3.75MG;3.75MG;3.75MG</u>	<u>A040440 005</u>	Oct 07, 2003
AB		<u>5MG;5MG;5MG;5MG</u>	<u>A040440 006</u>	Oct 07, 2003
AB		<u>7.5MG;7.5MG;7.5MG;7.5MG</u>	<u>A040440 007</u>	Oct 07, 2003
AB	SUN PHARM INDUSTRIES	<u>1.25MG;1.25MG;1.25MG;1.25MG</u>	<u>A040480 001</u>	Sep 09, 2003
AB		<u>1.875MG;1.875MG;1.875MG;1.875MG</u>	<u>A040480 002</u>	Sep 09, 2003
AB		<u>2.5MG;2.5MG;2.5MG;2.5MG</u>	<u>A040480 003</u>	Sep 09, 2003
AB		<u>3.125MG;3.125MG;3.125MG;3.125MG</u>	<u>A040480 004</u>	Sep 09, 2003

PRESCRIPTION DRUG PRODUCT LIST

AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATE

TABLET; ORAL

DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE

<u>AB</u>		<u>3.75MG; 3.75MG; 3.75MG; 3.75MG</u>	<u>A040480 005</u>	Sep 09, 2003
<u>AB</u>		<u>5MG; 5MG; 5MG; 5MG</u>	<u>A040480 006</u>	Sep 09, 2003
<u>AB</u>		<u>7.5MG; 7.5MG; 7.5MG; 7.5MG</u>	<u>A040480 007</u>	Sep 09, 2003
<u>AB</u>	SUNGEN PHARMA	<u>1.25MG; 1.25MG; 1.25MG; 1.25MG</u>	<u>A211352 001</u>	Dec 07, 2018
<u>AB</u>		<u>1.875MG; 1.875MG; 1.875MG; 1.875MG</u>	<u>A211352 002</u>	Dec 07, 2018
<u>AB</u>		<u>2.5MG; 2.5MG; 2.5MG; 2.5MG</u>	<u>A211352 003</u>	Dec 07, 2018
<u>AB</u>		<u>3.125MG; 3.125MG; 3.125MG; 3.125MG</u>	<u>A211352 004</u>	Dec 07, 2018
<u>AB</u>		<u>3.75MG; 3.75MG; 3.75MG; 3.75MG</u>	<u>A211352 005</u>	Dec 07, 2018
<u>AB</u>		<u>5MG; 5MG; 5MG; 5MG</u>	<u>A211352 006</u>	Dec 07, 2018
<u>AB</u>		<u>7.5MG; 7.5MG; 7.5MG; 7.5MG</u>	<u>A211352 007</u>	Dec 07, 2018

AMPHETAMINE SULFATE

TABLET; ORAL

AMPHETAMINE SULFATE

<u>AA</u>	AMNEAL PHARMS	<u>5MG</u>	<u>A211139 001</u>	Sep 26, 2018
<u>AA</u>		<u>10MG</u>	<u>A211139 002</u>	Sep 26, 2018
<u>AA</u>	AUROLIFE PHARMA LLC	<u>5MG</u>	<u>A211639 001</u>	Apr 17, 2019
<u>AA</u>		<u>10MG</u>	<u>A211639 002</u>	Apr 17, 2019
<u>AA</u>	BIONPHARMA INC	<u>5MG</u>	<u>A212919 001</u>	Nov 22, 2019
<u>AA</u>		<u>10MG</u>	<u>A212919 002</u>	Nov 22, 2019
<u>AA</u>	CEROVENE INC	<u>5MG</u>	<u>A212582 001</u>	Feb 04, 2020
<u>AA</u>		<u>10MG</u>	<u>A212582 002</u>	Feb 04, 2020
<u>AA</u>	GRANULES PHARMS	<u>5MG</u>	<u>A212619 001</u>	Aug 05, 2019
<u>AA</u>		<u>10MG</u>	<u>A212619 002</u>	Aug 05, 2019
<u>AA</u>	PRINSTON INC	<u>5MG</u>	<u>A211861 001</u>	Mar 11, 2020
<u>AA</u>		<u>10MG</u>	<u>A211861 002</u>	Mar 11, 2020

EVEKEO

<u>AA</u>	ARBOR PHARMS LLC	<u>5MG</u>	<u>A200166 001</u>	Aug 09, 2012
<u>AA</u>	!	<u>10MG</u>	<u>A200166 002</u>	Aug 09, 2012

TABLET, ORALLY DISINTEGRATING; ORAL

EVEKEO ODT

+	ARBOR PHARMS LLC	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

INJECTABLE; INJECTION

AMPHOTERICIN B

!	XGEN PHARMS	50MG/VIAL	A063206 001	Apr 29, 1992
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INJECTABLE, LIPID COMPLEX; INJECTION

ABELCET

+	LEADIANT BIOSCI INC	5MG/ML	N050724 001	Nov 20, 1995
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INJECTABLE, LIPOSOMAL; INJECTION

AMBISOME

+	ASTELLAS	50MG/VIAL	N050740 001	Aug 11, 1997
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AMPICILLIN SODIUM

INJECTABLE; INJECTION

AMPICILLIN SODIUM

<u>AP</u>	ACS DOBFAR SPA	<u>EQ 10GM BASE/VIAL</u>	<u>A090889 001</u>	Apr 03, 2013
<u>AP</u>	ANTIBIOTICE	<u>EQ 250MG BASE/VIAL</u>	<u>A090354 001</u>	Dec 28, 2009
<u>AP</u>		<u>EQ 500MG BASE/VIAL</u>	<u>A090354 002</u>	Dec 28, 2009
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>A090354 003</u>	Dec 28, 2009
<u>AP</u>		<u>EQ 2GM BASE/VIAL</u>	<u>A090354 004</u>	Dec 28, 2009
<u>AP</u>	AUROBINDO PHARMA	<u>EQ 250MG BASE/VIAL</u>	<u>A065499 002</u>	Aug 17, 2010
<u>AP</u>		<u>EQ 500MG BASE/VIAL</u>	<u>A065499 003</u>	Aug 17, 2010
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>A065499 004</u>	Aug 17, 2010
<u>AP</u>		<u>EQ 2GM BASE/VIAL</u>	<u>A065499 005</u>	Aug 17, 2010
<u>AP</u>		<u>EQ 10GM BASE/VIAL</u>	<u>A065493 001</u>	Aug 17, 2010
<u>AP</u>	HANFORD GC	<u>EQ 250MG BASE/VIAL</u>	<u>A062772 006</u>	Apr 15, 1993
<u>AP</u>		<u>EQ 500MG BASE/VIAL</u>	<u>A062772 007</u>	Apr 15, 1993
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>A062772 001</u>	Apr 15, 1993
<u>AP</u>		<u>EQ 2GM BASE/VIAL</u>	<u>A062772 003</u>	Apr 15, 1993
<u>AP</u>		<u>EQ 10GM BASE/VIAL</u>	<u>A063142 001</u>	Apr 15, 1993
<u>AP</u>	ISTITUTO BIO ITA SPA	<u>EQ 10GM BASE/VIAL</u>	<u>A201404 001</u>	Dec 20, 2013
<u>AP</u>		<u>EQ 250MG BASE/VIAL</u>	<u>A062719 001</u>	May 12, 1987
<u>AP</u>		<u>EQ 500MG BASE/VIAL</u>	<u>A062719 003</u>	May 12, 1987
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>A062719 002</u>	May 12, 1987

PRESCRIPTION DRUG PRODUCT LIST

AMPICILLIN SODIUM

INJECTABLE; INJECTION

AMPICILLIN SODIUM

AP		<u>EQ 2GM BASE/VIAL</u>	<u>A062797 002</u>	Jul 12, 1993
AP	MYLAN LABS LTD	<u>EQ 250MG BASE/VIAL</u>	<u>A201025 001</u>	Apr 09, 2014
AP		<u>EQ 500MG BASE/VIAL</u>	<u>A201025 002</u>	Apr 09, 2014
AP		<u>EQ 1GM BASE/VIAL</u>	<u>A201025 003</u>	Apr 09, 2014
AP		<u>EQ 2GM BASE/VIAL</u>	<u>A201025 004</u>	Apr 09, 2014
AP		<u>EQ 10GM BASE/VIAL</u>	<u>A202198 001</u>	Apr 07, 2014
AP	SAGENT PHARMS INC	<u>EQ 125MG BASE/VIAL</u>	<u>A090583 001</u>	Nov 27, 2015
AP		<u>EQ 250MG BASE/VIAL</u>	<u>A090583 002</u>	Nov 27, 2015
AP		<u>EQ 500MG BASE/VIAL</u>	<u>A090583 003</u>	Nov 27, 2015
AP		<u>EQ 1GM BASE/VIAL</u>	<u>A090583 004</u>	Nov 27, 2015
AP		<u>EQ 2GM BASE/VIAL</u>	<u>A090583 005</u>	Nov 27, 2015
AP		<u>EQ 10GM BASE/VIAL</u>	<u>A090581 001</u>	Oct 20, 2015
AP	! SANDOZ	<u>EQ 125MG BASE/VIAL</u>	<u>A061395 001</u>	
AP	!	<u>EQ 250MG BASE/VIAL</u>	<u>A061395 002</u>	
AP	!	<u>EQ 500MG BASE/VIAL</u>	<u>A061395 003</u>	
AP	!	<u>EQ 1GM BASE/VIAL</u>	<u>A061395 004</u>	
AP	!	<u>EQ 2GM BASE/VIAL</u>	<u>A061395 005</u>	
AP	!	<u>EQ 10GM BASE/VIAL</u>	<u>A061395 006</u>	

POWDER; INTRAVENOUS

AMPICILLIN SODIUM

AP	! SANDOZ	<u>EQ 1GM BASE/VIAL</u>	<u>A062738 001</u>	Feb 19, 1987
AP	!	<u>EQ 2GM BASE/VIAL</u>	<u>A062738 002</u>	Feb 19, 1987

AMPICILLIN SODIUM; SULBACTAM SODIUM

INJECTABLE; INJECTION

AMPICILLIN AND SULBACTAM

AP	ACS DOBFAR	<u>EQ 1GM BASE/VIAL;EQ 500MG BASE/VIAL</u>	<u>A065406 001</u>	Dec 22, 2009
AP		<u>EQ 2GM BASE/VIAL;EQ 1GM BASE/VIAL</u>	<u>A065406 002</u>	Dec 22, 2009
AP		<u>EQ 10GM BASE/VIAL;EQ 5GM BASE/VIAL</u>	<u>A065403 001</u>	Dec 23, 2009
AP	ANTIBIOTICE	<u>EQ 1GM BASE/VIAL;EQ 500MG BASE/VIAL</u>	<u>A201406 001</u>	Dec 07, 2015
AP		<u>EQ 2GM BASE/VIAL;EQ 1GM BASE/VIAL</u>	<u>A201406 002</u>	Dec 07, 2015
AP	ASTRAL	<u>EQ 1GM BASE/VIAL;EQ 500MG BASE/VIAL</u>	<u>A090579 001</u>	Jan 08, 2016
AP		<u>EQ 2GM BASE/VIAL;EQ 1GM BASE/VIAL</u>	<u>A090579 002</u>	Jan 08, 2016
AP		<u>EQ 10GM BASE/VIAL;EQ 5GM BASE/VIAL</u>	<u>A090578 001</u>	Jan 11, 2016
AP	AUROBINDO PHARMA	<u>EQ 1GM BASE/VIAL;EQ 500MG BASE/VIAL</u>	<u>A090340 001</u>	Sep 20, 2010
AP		<u>EQ 1GM BASE/VIAL;EQ 500MG BASE/VIAL</u>	<u>A090349 001</u>	Sep 20, 2010
AP		<u>EQ 2GM BASE/VIAL;EQ 1GM BASE/VIAL</u>	<u>A090340 002</u>	Sep 20, 2010
AP		<u>EQ 2GM BASE/VIAL;EQ 1GM BASE/VIAL</u>	<u>A090349 002</u>	Sep 20, 2010
AP		<u>EQ 10GM BASE/VIAL;EQ 5GM BASE/VIAL</u>	<u>A090339 001</u>	Sep 20, 2010
AP	HANFORD GC	<u>EQ 1GM BASE/VIAL;EQ 500MG BASE/VIAL</u>	<u>A065176 001</u>	Nov 30, 2005
AP		<u>EQ 2GM BASE/VIAL;EQ 1GM BASE/VIAL</u>	<u>A065176 002</u>	Nov 30, 2005
AP		<u>EQ 10GM BASE/VIAL;EQ 5GM BASE/VIAL</u>	<u>A065188 001</u>	Nov 25, 2005
AP	ISTITUTO BIO ITA SPA	<u>EQ 1GM BASE/VIAL;EQ 500MG BASE/VIAL</u>	<u>A065222 001</u>	Nov 29, 2005
AP		<u>EQ 2GM BASE/VIAL;EQ 1GM BASE/VIAL</u>	<u>A065222 002</u>	Nov 29, 2005
AP		<u>EQ 10GM BASE/VIAL;EQ 5GM BASE/VIAL</u>	<u>A065314 001</u>	Nov 27, 2006
AP	MYLAN LABS LTD	<u>EQ 1GM BASE/VIAL;EQ 500MG BASE/VIAL</u>	<u>A201024 001</u>	Apr 07, 2014
AP		<u>EQ 2GM BASE/VIAL;EQ 1GM BASE/VIAL</u>	<u>A201024 002</u>	Apr 07, 2014
AP		<u>EQ 10GM BASE/VIAL;EQ 5GM BASE/VIAL</u>	<u>A202197 001</u>	Apr 07, 2014
AP	SANDOZ	<u>EQ 1GM BASE/VIAL;EQ 500MG BASE/VIAL</u>	<u>A065241 001</u>	Jul 25, 2006
AP		<u>EQ 1GM BASE/VIAL;EQ 500MG BASE/VIAL</u>	<u>A065310 001</u>	Jul 25, 2006
AP		<u>EQ 2GM BASE/VIAL;EQ 1GM BASE/VIAL</u>	<u>A065241 002</u>	Jul 25, 2006
AP		<u>EQ 2GM BASE/VIAL;EQ 1GM BASE/VIAL</u>	<u>A065310 002</u>	Jul 25, 2006
AP		<u>EQ 10GM BASE/VIAL;EQ 5GM BASE/VIAL</u>	<u>A065240 001</u>	Jul 25, 2006
AP	WEST-WARD PHARMS INT	<u>EQ 1GM BASE/VIAL;EQ 500MG BASE/VIAL</u>	<u>A065074 001</u>	Mar 19, 2002
AP		<u>EQ 2GM BASE/VIAL;EQ 1GM BASE/VIAL</u>	<u>A065074 002</u>	Mar 19, 2002
AP		<u>EQ 10GM BASE/VIAL;EQ 5GM BASE/VIAL</u>	<u>A065076 001</u>	Mar 19, 2002
<u>UNASYN</u>				
AP	! PFIZER	<u>EQ 2GM BASE/VIAL;EQ 1GM BASE/VIAL</u>	<u>A062901 002</u>	Feb 27, 1992
AP	!	<u>EQ 1GM BASE/VIAL;EQ 500MG BASE/VIAL</u>	<u>A062901 001</u>	Nov 23, 1988
AP	+!	<u>EQ 1GM BASE/VIAL;EQ 500MG BASE/VIAL</u>	<u>N050608 002</u>	Dec 31, 1986
AP	+!	<u>EQ 2GM BASE/VIAL;EQ 1GM BASE/VIAL</u>	<u>N050608 001</u>	Dec 31, 1986
AP	+!	<u>EQ 10GM BASE/VIAL;EQ 5GM BASE/VIAL</u>	<u>N050608 005</u>	Dec 10, 1993

PRESCRIPTION DRUG PRODUCT LIST

AMPICILLIN/AMPICILLIN TRIHYDRATE

CAPSULE; ORAL

AMPICILLIN TRIHYDRATE

<u>AB</u>	DAVA PHARMS INC	<u>EQ 250MG BASE</u>	<u>A062883 001</u>	Feb 25, 1988
<u>AB</u>	!	<u>EQ 500MG BASE</u>	<u>A062882 001</u>	Feb 25, 1988
<u>AB</u>	SANDOZ	<u>EQ 250MG BASE</u>	<u>A064082 001</u>	Aug 29, 1995
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>A064082 002</u>	Aug 29, 1995
FOR SUSPENSION; ORAL				
AMPICILLIN TRIHYDRATE				
	DAVA PHARMS INC	EQ 125MG BASE/5ML	A062982 001	Feb 10, 1989
	!	EQ 250MG BASE/5ML	A062982 002	Feb 10, 1989

ANAGRELIDE HYDROCHLORIDE

CAPSULE; ORAL

AGRYLIN

<u>AB</u>	SHIRE LLC	<u>EQ 0.5MG BASE</u>	<u>N020333 001</u>	Mar 14, 1997
<u>ANAGRELIDE HYDROCHLORIDE</u>				
<u>AB</u>	BARR	<u>EQ 0.5MG BASE</u>	<u>A076530 001</u>	Apr 18, 2005
<u>AB</u>		<u>EQ 1MG BASE</u>	<u>A076530 002</u>	Apr 18, 2005
<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>	FRESENIUS KABI USA	<u>1MG</u>	<u>A090088 001</u>	Jun 28, 2010
<u>AB</u>	HIKMA	<u>1MG</u>	<u>A078485 001</u>	Jun 28, 2010
<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>	NEOPHARMA	<u>1MG</u>	<u>A090732 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

ARIMIDEX

<u>AB</u>	+!	ANI PHARMS INC	<u>1MG</u>	<u>N020541 001</u>	Dec 27, 1995
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ANGIOTENSIN II ACETATE

SOLUTION; INTRAVENOUS

GIAPREZA

	+!	LA JOLLA PHARMA	EQ 2.5MG BASE/ML (EQ 2.5MG BASE/ML)	N209360 001	Dec 21, 2017
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ANIDULAFUNGIN

POWDER; INTRAVENOUS

ERAXIS

	+!	VICURON	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
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APIXABAN

TABLET; ORAL

APIXABAN

<u>AB</u>	MICRO LABS	<u>2.5MG</u>	<u>A210013 001</u>	Dec 23, 2019
<u>AB</u>		<u>5MG</u>	<u>A210013 002</u>	Dec 23, 2019

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

PRESCRIPTION DRUG PRODUCT LIST

APOMORPHINE HYDROCHLORIDE

INJECTABLE; SUBCUTANEOUS

APOKYN

+! US WORLDMEDS 30MG/3ML (10MG/ML) N021264 002 Apr 20, 2004

APRACLONIDINE HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

APRACLONIDINE HYDROCHLORIDE**AT** AKORN INC **EQ 0.5% BASE** **A077764 001** Mar 12, 2009IOPIDINE**AT** +! NOVARTIS **EQ 0.5% BASE** **N020258 001** Jul 30, 1993

+! EQ 1% BASE N019779 001 Dec 31, 1987

APREMLLAST

TABLET; ORAL

OTEZLA

+ AMGEN INC 10MG N205437 001 Mar 21, 2014

+ 20MG N205437 002 Mar 21, 2014

+! 30MG N205437 003 Mar 21, 2014

APREPITANT

CAPSULE; ORAL

APREPITANT**AB** GLENMARK PHARMS SA **40MG** **A207777 001** Oct 12, 2017**AB** **80MG** **A207777 002** Oct 12, 2017**AB** **125MG** **A207777 003** Oct 12, 2017**AB** SANDOZ **40MG** **A090999 001** Sep 24, 2012**AB** **80MG** **A090999 002** Sep 24, 2012**AB** **125MG** **A090999 003** Sep 24, 2012EMEND**AB** + MERCK **80MG** **N021549 001** Mar 26, 2003**AB** +! **125MG** **N021549 002** Mar 26, 2003

EMULSION; INTRAVENOUS

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

ARFORMOTEROL TARTRATE

SOLUTION; INHALATION

BROVANA

+! SUNOVION EQ 0.015MG BASE/2ML N021912 001 Oct 06, 2006

ARGATROBAN

INJECTABLE; INJECTION

ARGATROBAN**AP** AMNEAL PHARMS CO **250MG/2.5ML (100MG/ML)** **A206698 001** Jan 26, 2018**AP** FRESENIUS KABI USA **250MG/2.5ML (100MG/ML)** **N201811 001** Mar 23, 2015**AP** HIKMA PHARM CO LTD **250MG/2.5ML (100MG/ML)** **N203049 001** Jan 05, 2012**AP** HOSPIRA INC **250MG/2.5ML (100MG/ML)** **A204120 001** Sep 21, 2016**AP** MYLAN INSTITUTIONAL **250MG/2.5ML (100MG/ML)** **A202626 001** Jun 30, 2014**AP** +! NOVARTIS **250MG/2.5ML (100MG/ML)** **N020883 001** Jun 30, 2000**AP** PAR STERILE **250MG/2.5ML (100MG/ML)** **A091665 001** Jun 30, 2014

PRODUCTS

+! HIKMA PHARM CO LTD 50MG/50ML (1MG/ML) N203049 002 Sep 30, 2016

INJECTABLE; INTRAVENOUS

ARGATROBAN IN SODIUM CHLORIDE**AP** GLAND PHARMA LTD **125MG/125ML (1MG/ML)** **A205570 001** May 22, 2017**AP** +! SANDOZ **125MG/125ML (1MG/ML)** **N022485 001** May 09, 2011

ARGATROBAN IN 0.9% SODIUM CHLORIDE

TEVA PHARMS USA 250MG/250ML (1MG/ML) N206769 001 Dec 15, 2014

ARGATROBAN IN SODIUM CHLORIDE

+! EAGLE PHARMS 50MG/50ML (1MG/ML) N022434 001 Jun 29, 2011

SOLUTION; INTRAVENOUS

ARGATROBAN IN SODIUM CHLORIDE

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

LTD

ARGININE HYDROCHLORIDE

INJECTABLE; INJECTION

R-GENE 10

+! PHARMACIA AND 10GM/100ML N016931 001

UPJOHN

PRESCRIPTION DRUG PRODUCT LIST

ARIPIPIRAZOLE

FOR SUSPENSION, EXTENDED RELEASE; INTRAMUSCULAR

ABILIFY MAINTENA KIT

+	OTSUKA PHARM CO LTD	300MG/VIAL	N202971	001	Feb 28, 2013
+		300MG	N202971	003	Sep 29, 2014
+	!	400MG/VIAL	N202971	002	Feb 28, 2013
+		400MG	N202971	004	Sep 29, 2014

SOLUTION; ORAL

ARIPIPIRAZOLE

AA	!	AMNEAL PHARMS	<u>1MG/ML</u>	<u>A203906</u>	<u>001</u>	Aug 14, 2015
AA		APOTEX INC	<u>1MG/ML</u>	<u>A204094</u>	<u>001</u>	Sep 30, 2015
AA		AUROBINDO PHARMA LTD	<u>1MG/ML</u>	<u>A210479</u>	<u>001</u>	Jan 29, 2019
AA		LANNETT CO INC	<u>1MG/ML</u>	<u>A204171</u>	<u>001</u>	Aug 14, 2015
AA		VISTAPHARM	<u>1MG/ML</u>	<u>A212870</u>	<u>001</u>	Dec 26, 2019

TABLET; ORAL

ABILIFY

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

ARIPIPIRAZOLE

AB		ACCORD HLTHCARE	<u>2MG</u>	<u>A206251</u>	<u>001</u>	Dec 07, 2016
AB			<u>5MG</u>	<u>A206251</u>	<u>002</u>	Dec 07, 2016
AB			<u>10MG</u>	<u>A206251</u>	<u>003</u>	Dec 07, 2016
AB			<u>15MG</u>	<u>A206251</u>	<u>004</u>	Dec 07, 2016
AB			<u>20MG</u>	<u>A206251</u>	<u>005</u>	Dec 07, 2016
AB			<u>30MG</u>	<u>A206251</u>	<u>006</u>	Dec 07, 2016
AB		AJANTA PHARMA LTD	<u>2MG</u>	<u>A206174</u>	<u>001</u>	Sep 12, 2016
AB			<u>5MG</u>	<u>A206174</u>	<u>002</u>	Sep 12, 2016
AB			<u>10MG</u>	<u>A206174</u>	<u>003</u>	Sep 12, 2016
AB			<u>15MG</u>	<u>A206174</u>	<u>004</u>	Sep 12, 2016
AB			<u>20MG</u>	<u>A206174</u>	<u>005</u>	Sep 12, 2016
AB			<u>30MG</u>	<u>A206174</u>	<u>006</u>	Sep 12, 2016
AB		ALEMBIC PHARMS LTD	<u>2MG</u>	<u>A202101</u>	<u>001</u>	Apr 28, 2015
AB			<u>5MG</u>	<u>A202101</u>	<u>002</u>	Apr 28, 2015
AB			<u>10MG</u>	<u>A202101</u>	<u>003</u>	Apr 28, 2015
AB			<u>15MG</u>	<u>A202101</u>	<u>004</u>	Apr 28, 2015
AB			<u>20MG</u>	<u>A202101</u>	<u>005</u>	Apr 28, 2015
AB			<u>30MG</u>	<u>A202101</u>	<u>006</u>	Apr 28, 2015
AB		ALKEM LABS LTD	<u>2MG</u>	<u>A207105</u>	<u>001</u>	Feb 21, 2019
AB			<u>5MG</u>	<u>A207105</u>	<u>002</u>	Feb 21, 2019
AB			<u>10MG</u>	<u>A207105</u>	<u>003</u>	Feb 21, 2019
AB			<u>15MG</u>	<u>A207105</u>	<u>004</u>	Feb 21, 2019
AB			<u>20MG</u>	<u>A207105</u>	<u>005</u>	Feb 21, 2019
AB			<u>30MG</u>	<u>A207105</u>	<u>006</u>	Feb 21, 2019
AB		AMNEAL PHARMS	<u>2MG</u>	<u>A204838</u>	<u>001</u>	Jun 17, 2016
AB			<u>5MG</u>	<u>A204838</u>	<u>002</u>	Jun 17, 2016
AB			<u>10MG</u>	<u>A204838</u>	<u>003</u>	Jun 17, 2016
AB			<u>15MG</u>	<u>A204838</u>	<u>004</u>	Jun 17, 2016
AB			<u>20MG</u>	<u>A204838</u>	<u>005</u>	Jun 17, 2016
AB			<u>30MG</u>	<u>A204838</u>	<u>006</u>	Jun 17, 2016
AB		APOTEX INC	<u>2MG</u>	<u>A078583</u>	<u>001</u>	Jul 24, 2015
AB			<u>5MG</u>	<u>A078583</u>	<u>002</u>	Jul 24, 2015
AB			<u>10MG</u>	<u>A078583</u>	<u>003</u>	Jul 24, 2015
AB			<u>15MG</u>	<u>A078583</u>	<u>004</u>	Jul 24, 2015
AB			<u>20MG</u>	<u>A078583</u>	<u>005</u>	Jul 24, 2015
AB			<u>30MG</u>	<u>A078583</u>	<u>006</u>	Jul 24, 2015
AB		AUROBINDO PHARMA LTD	<u>2MG</u>	<u>A203908</u>	<u>001</u>	Oct 08, 2015
AB			<u>5MG</u>	<u>A203908</u>	<u>002</u>	Oct 08, 2015
AB			<u>10MG</u>	<u>A203908</u>	<u>003</u>	Oct 08, 2015
AB			<u>15MG</u>	<u>A203908</u>	<u>004</u>	Oct 08, 2015
AB			<u>20MG</u>	<u>A203908</u>	<u>005</u>	Oct 08, 2015
AB			<u>30MG</u>	<u>A203908</u>	<u>006</u>	Oct 08, 2015
AB		BOSCOGEN	<u>2MG</u>	<u>A091279</u>	<u>001</u>	Jan 09, 2017
AB			<u>5MG</u>	<u>A091279</u>	<u>002</u>	Jan 09, 2017
AB			<u>10MG</u>	<u>A091279</u>	<u>003</u>	Jan 09, 2017
AB			<u>15MG</u>	<u>A091279</u>	<u>004</u>	Jan 09, 2017
AB			<u>20MG</u>	<u>A091279</u>	<u>005</u>	Jan 09, 2017
AB			<u>30MG</u>	<u>A091279</u>	<u>006</u>	Jan 09, 2017

PRESCRIPTION DRUG PRODUCT LIST

ARIPIPIRAZOLE

TABLET;ORAL

ARIPIPIRAZOLE

<u>AB</u>	HETERO LABS LTD V	<u>2MG</u>	<u>A205064 001</u>	Apr 28, 2015
<u>AB</u>		<u>5MG</u>	<u>A205064 002</u>	Apr 28, 2015
<u>AB</u>		<u>10MG</u>	<u>A205064 003</u>	Apr 28, 2015
<u>AB</u>		<u>15MG</u>	<u>A205064 004</u>	Apr 28, 2015
<u>AB</u>		<u>20MG</u>	<u>A205064 005</u>	Apr 28, 2015
<u>AB</u>		<u>30MG</u>	<u>A205064 006</u>	Apr 28, 2015
<u>AB</u>	MACLEODS PHARMS LTD	<u>2MG</u>	<u>A204111 001</u>	Oct 07, 2016
<u>AB</u>		<u>5MG</u>	<u>A204111 002</u>	Oct 07, 2016
<u>AB</u>		<u>10MG</u>	<u>A204111 003</u>	Oct 07, 2016
<u>AB</u>		<u>15MG</u>	<u>A204111 004</u>	Oct 07, 2016
<u>AB</u>		<u>20MG</u>	<u>A204111 005</u>	Oct 07, 2016
<u>AB</u>		<u>30MG</u>	<u>A204111 006</u>	Oct 07, 2016
<u>AB</u>	ORCHID HLTHCARE	<u>2MG</u>	<u>A202683 001</u>	May 23, 2017
<u>AB</u>		<u>5MG</u>	<u>A202683 002</u>	May 23, 2017
<u>AB</u>		<u>10MG</u>	<u>A202683 003</u>	May 23, 2017
<u>AB</u>		<u>15MG</u>	<u>A202683 004</u>	May 23, 2017
<u>AB</u>		<u>20MG</u>	<u>A202683 005</u>	May 23, 2017
<u>AB</u>		<u>30MG</u>	<u>A202683 006</u>	May 23, 2017
<u>AB</u>	PRINSTON INC	<u>2MG</u>	<u>A205363 001</u>	Dec 04, 2017
<u>AB</u>		<u>5MG</u>	<u>A205363 002</u>	Dec 04, 2017
<u>AB</u>		<u>10MG</u>	<u>A205363 003</u>	Dec 04, 2017
<u>AB</u>		<u>15MG</u>	<u>A205363 004</u>	Dec 04, 2017
<u>AB</u>		<u>20MG</u>	<u>A205363 005</u>	Dec 04, 2017
<u>AB</u>		<u>30MG</u>	<u>A205363 006</u>	Dec 04, 2017
<u>AB</u>	SCIEGEN PHARMS INC	<u>2MG</u>	<u>A206383 001</u>	Sep 29, 2016
<u>AB</u>		<u>5MG</u>	<u>A206383 002</u>	Sep 29, 2016
<u>AB</u>		<u>10MG</u>	<u>A206383 003</u>	Sep 29, 2016
<u>AB</u>		<u>15MG</u>	<u>A206383 004</u>	Sep 29, 2016
<u>AB</u>		<u>20MG</u>	<u>A206383 005</u>	Sep 29, 2016
<u>AB</u>		<u>30MG</u>	<u>A206383 006</u>	Sep 29, 2016
<u>AB</u>	TORRENT	<u>2MG</u>	<u>A201519 001</u>	Apr 28, 2015
<u>AB</u>		<u>10MG</u>	<u>A201519 003</u>	Apr 28, 2015
<u>AB</u>		<u>5MG</u>	<u>A201519 002</u>	Apr 28, 2015
<u>AB</u>		<u>15MG</u>	<u>A201519 004</u>	Apr 28, 2015
<u>AB</u>		<u>20MG</u>	<u>A201519 005</u>	Apr 28, 2015
<u>AB</u>		<u>30MG</u>	<u>A201519 006</u>	Apr 28, 2015
	ABILIFY MYCITE KIT			
	+ OTSUKA PHARM CO LTD	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

TABLET, ORALLY DISINTEGRATING;ORAL

ARIPIPIRAZOLE

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

ARIPIPIRAZOLE 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	N209830 001	Jun 29, 2018

PRESCRIPTION DRUG PRODUCT LIST

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>	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>	INGENUS PHARMS LLC	<u>1MG/ML</u>	<u>A209315 001</u>	Nov 15, 2018
<u>AP</u>	NEXUS PHARMS	<u>1MG/ML</u>	<u>A209780 001</u>	Nov 15, 2018
<u>AP</u>	STI PHARMA LLC	<u>1MG/ML</u>	<u>A209873 001</u>	May 06, 2019
<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
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ARTEMETHER; LUMEFANTRINE

TABLET; ORAL

COARTEM

+!	NOVARTIS	20MG;120MG	N022268 001	Apr 07, 2009
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ARTICAINE HYDROCHLORIDE; EPINEPHRINE BITARTRATE

INJECTABLE; INJECTION

SEPTOCAINE

<u>AP</u>	+! DEPROCO	<u>4%;EQ 0.0085MG BASE/1.7ML (4%;EQ 0.005MG BASE/ML)</u>	<u>N022010 001</u>	Mar 30, 2006
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ULTACAN

<u>AP</u>	HANSAMED INC	<u>4%;EQ 0.0085MG BASE/1.7ML (4%; EQ 0.005MG BASE/ML)</u>	<u>A201751 001</u>	Jul 11, 2017
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ORABLOC

+	PIERREL	4%;EQ 0.009MG BASE/1.8ML (EQ 0.005MG BASE/ML)	N022466 001	Feb 26, 2010
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+!		4%;EQ 0.018MG BASE/1.8ML (EQ 0.01MG BASE/ML)	N022466 002	Feb 26, 2010
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SEPTOCAINE

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

SOLUTION; INTRAVENOUS

ASCOR

+!	MCGUFF	25,000MG/50ML (500MG/ML)	N209112 001	Oct 02, 2017
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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 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	N021265 001	Feb 21, 2001
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PRESCRIPTION DRUG PRODUCT LIST

3-41 (of 453)

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

INJECTABLE; INTRAVENOUS

INFUVITE PEDIATRIC (PHARMACY BULK PACKAGE)

+	!	SANDOZ CANADA INC	80MG/VIAL; 0.02MG/VIAL; 400 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	N021265 002	Jan 29, 2004
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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; 17MG/VIAL; 0.2MG/VIAL; 1MG/VIAL; 1.4MG/VIAL; EQ 1.2MG BASE/VIAL; 0.7MG/VIAL; 7MG/VIAL	N018920 001	Sep 21, 2000
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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; 15MG/VIAL; 0.005MG/VIAL; 0.6MG/VIAL; 40MG/VIAL; 6MG/VIAL; 3.6MG/VIAL; 6MG/VIAL; 1MG/VIAL; 10MG/VIAL; 0.15MG/VIAL	N021625 001	Jan 30, 2004
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M.V.I. ADULT (PHARMACY BULK PACKAGE)

+	!	HOSPIRA	200MG/5ML; 0.06MG/5ML; 0.005MG/5ML; 15MG/5ML; 0.005MG/5ML; 0.6MG/5ML; 40MG/5ML; 6MG/5ML; 3.6MG/5ML; 6MG/5ML; 1MG/5ML; 10MG/5ML; 0.15MG/5ML	N021643 001	Feb 18, 2004
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ASCORBIC ACID; POLYETHYLENE GLYCOL 3350; POTASSIUM CHLORIDE; SODIUM ASCORBATE; SODIUM CHLORIDE; SODIUM SULFATE

FOR SOLUTION; ORAL

MOVIPREP

AA	+	!	SALIX PHARMS	4.7GM; 100GM; 1.015GM; 5.9GM; 2.691GM; 7.5GM	N021881 001	Aug 02, 2006
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PEG-3350, SODIUM SULFATE, SODIUM CHLORIDE, POTASSIUM CHLORIDE, SODIUM ASCORBATE AND ASCORBIC

AA			NOVEL LABS INC	4.7GM; 100GM; 1.015GM; 5.9GM; 2.691GM; 7.5GM	A090145 001	Jan 25, 2012
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PLENVU

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

SAPHRIS

+		ALLERGAN	EQ 2.5MG BASE	N022117 003	Mar 12, 2015
+			EQ 5MG BASE	N022117 001	Aug 13, 2009
+	!		EQ 10MG BASE	N022117 002	Aug 13, 2009

ASPIRIN

CAPSULE, EXTENDED RELEASE; ORAL

DURLAZA

+	!	ESPERO	162.5MG	N200671 001	Sep 04, 2015
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ASPIRIN; BUTALBITAL; CAFFEINE

CAPSULE; ORAL

FIORINAL

AA	+	!	ALLERGAN	325MG; 50MG; 40MG	N017534 005	Apr 16, 1986
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LANORINAL

AA			LANNETT	325MG; 50MG; 40MG	A086996 002	Oct 11, 1985
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TABLET; ORAL

BUTALBITAL, ASPIRIN AND CAFFEINE

AA	!		HIKMA INTL PHARMS	325MG; 50MG; 40MG	A086162 002	Feb 16, 1984
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AA			PII	325MG; 50MG; 40MG	A204195 001	Sep 22, 2016
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PRESCRIPTION DRUG PRODUCT LIST

ASPIRIN; BUTALBITAL; CAFFEINE; CODEINE PHOSPHATE

CAPSULE; ORAL

BUTALBITAL, ASPIRIN, CAFFEINE, AND CODEINE PHOSPHATE

AB	MAYNE PHARMA INC	<u>325MG; 50MG; 40MG; 30MG</u>	<u>A203335 001</u>	Oct 30, 2015
AB	NEXGEN PHARMA INC	<u>325MG; 50MG; 40MG; 30MG</u>	<u>A075231 001</u>	Nov 30, 2001
AB	STEVENS J	<u>325MG; 50MG; 40MG; 30MG</u>	<u>A074951 001</u>	Aug 31, 1998

FIORINAL W/CODEINE

AB	+ ! ALLERGAN	<u>325MG; 50MG; 40MG; 30MG</u>	<u>N019429 003</u>	Oct 26, 1990
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ASPIRIN; CARISOPRODOL

TABLET; ORAL

CARISOPRODOL AND ASPIRIN

AB	! HERITAGE PHARMS INC	<u>325MG; 200MG</u>	<u>A089594 001</u>	Mar 31, 1989
AB	NOVAST LABS	<u>325MG; 200MG</u>	<u>A040832 001</u>	Jan 07, 2010
AB	SANDOZ	<u>325MG; 200MG</u>	<u>A040116 001</u>	Apr 25, 1996

ASPIRIN; CARISOPRODOL; CODEINE PHOSPHATE

TABLET; ORAL

CARISOPRODOL, ASPIRIN AND CODEINE PHOSPHATE

AB	INGENUS PHARMS NJ	<u>325MG; 200MG; 16MG</u>	<u>A040860 001</u>	Jan 07, 2010
AB	! SANDOZ	<u>325MG; 200MG; 16MG</u>	<u>A040118 001</u>	Apr 16, 1996

ASPIRIN; DIPYRIDAMOLE

CAPSULE, EXTENDED RELEASE; ORAL

AGGRENOL

AB	+ ! BOEHRINGER INGELHEIM	<u>25MG; 200MG</u>	<u>N020884 001</u>	Nov 22, 1999
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ASPIRIN AND DIPYRIDAMOLE

AB	AMNEAL PHARMS	<u>25MG; 200MG</u>	<u>A206392 001</u>	Mar 08, 2016
AB	BARR	<u>25MG; 200MG</u>	<u>A078804 001</u>	Aug 14, 2009
AB	DR REDDYS	<u>25MG; 200MG</u>	<u>A209048 001</u>	Oct 10, 2018
AB	GLENMARK PHARMS SA	<u>25MG; 200MG</u>	<u>A210318 001</u>	May 24, 2019
AB	LANNETT CO INC	<u>25MG; 200MG</u>	<u>A204552 001</u>	Mar 20, 2019
AB	PAR PHARM INC	<u>25MG; 200MG</u>	<u>A207944 001</u>	Jan 18, 2017
AB	SANDOZ INC	<u>25MG; 200MG</u>	<u>A206739 001</u>	Jan 18, 2017
AB	SUN PHARM	<u>25MG; 200MG</u>	<u>A208572 001</u>	Aug 21, 2018
AB	ZYDUS PHARMS	<u>25MG; 200MG</u>	<u>A206753 001</u>	Aug 29, 2017

ASPIRIN; METHOCARBAMOL

TABLET; ORAL

METHOCARBAMOL AND ASPIRIN

!	STEVENS J	325MG; 400MG	A081145 001	Jan 31, 1995
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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

AA	ACTAVIS LABS FL INC	<u>325MG; 4.8355MG</u>	<u>A090084 001</u>	Mar 22, 2011
AA	MAYNE PHARMA INC	<u>325MG; 4.8355MG</u>	<u>A091670 001</u>	Mar 16, 2011

PERCODAN

AA	+ ! ENDO PHARMS	<u>325MG; 4.8355MG</u>	<u>N007337 007</u>	Aug 05, 2005
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ATAZANAVIR SULFATE

CAPSULE; ORAL

ATAZANAVIR SULFATE

AB	AUROBINDO PHARMA LTD	<u>EQ 100MG BASE</u>	<u>A204806 001</u>	Jun 25, 2018
AB		<u>EQ 150MG BASE</u>	<u>A204806 002</u>	Jun 25, 2018
AB		<u>EQ 200MG BASE</u>	<u>A204806 003</u>	Jun 25, 2018
AB		<u>EQ 300MG BASE</u>	<u>A204806 004</u>	Jun 25, 2018
AB	CIPLA	<u>EQ 100MG BASE</u>	<u>A200626 001</u>	Aug 09, 2018
AB		<u>EQ 150MG BASE</u>	<u>A200626 002</u>	Aug 09, 2018
AB		<u>EQ 200MG BASE</u>	<u>A200626 003</u>	Aug 09, 2018
AB		<u>EQ 300MG BASE</u>	<u>A200626 004</u>	Aug 09, 2018
AB	TEVA PHARMS USA	<u>EQ 100MG BASE</u>	<u>A091673 001</u>	Apr 22, 2014
AB		<u>EQ 150MG BASE</u>	<u>A091673 002</u>	Apr 22, 2014
AB		<u>EQ 200MG BASE</u>	<u>A091673 003</u>	Apr 22, 2014
AB		<u>EQ 300MG BASE</u>	<u>A091673 004</u>	Apr 22, 2014

REYATAZ

AB	+ BRISTOL MYERS SQUIBB	<u>EQ 150MG BASE</u>	<u>N021567 002</u>	Jun 20, 2003
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PRESCRIPTION DRUG PRODUCT LIST

ATAZANAVIR SULFATE

CAPSULE;ORAL

REYATAZ

AB + EQ 200MG BASE N021567 003 Jun 20, 2003
AB +! EQ 300MG BASE N021567 004 Oct 16, 2006

POWDER;ORAL

REYATAZ

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

ATAZANAVIR SULFATE; COBICISTAT

TABLET;ORAL

EVOTAZ

+! BRISTOL-MYERS EQ 300MG BASE;150MG N206353 001 Jan 29, 2015
 SQUIBB

ATENOLOL

TABLET;ORAL

ATENOLOL

AB ALVOGEN 25MG A072304 002 Jul 31, 1992
AB 50MG A072304 003 Jul 18, 1988
AB 100MG A072304 001 Jul 15, 1988
AB AUROBINDO PHARMA 25MG A078512 001 Oct 31, 2007
AB 50MG A078512 002 Oct 31, 2007
AB 100MG A078512 003 Oct 31, 2007
AB DAVA PHARMS INC 50MG A073542 001 Dec 19, 1991
AB 100MG A073543 001 Dec 19, 1991
AB HLTHCARE 25MG A074052 001 May 01, 1992
AB 50MG A073025 001 Sep 17, 1991
AB 100MG A073026 001 Sep 17, 1991
AB IPCA LABS LTD 25MG A077877 001 Dec 27, 2006
AB 50MG A077877 002 Dec 27, 2006
AB 100MG A077877 003 Dec 27, 2006
AB MYLAN 25MG A073457 002 Apr 26, 1999
AB 50MG A073457 003 Jan 24, 1992
AB 100MG A073457 001 Jan 24, 1992
AB TEVA 25MG A074056 003 Jul 19, 2004
AB 50MG A074056 001 Jan 18, 1995
AB 100MG A074056 002 Jan 18, 1995
AB UNICHEM LABS LTD 25MG A213136 001 Nov 21, 2019
AB 50MG A213136 002 Nov 21, 2019
AB 100MG A213136 003 Nov 21, 2019
AB UNIQUE PHARM LABS 25MG A077443 001 Sep 13, 2006
AB 50MG A077443 002 Sep 13, 2006
AB 100MG A077443 003 Sep 13, 2006
AB ZYDUS PHARMS USA 25MG A076900 001 Jan 28, 2005
AB 50MG A076900 002 Jan 28, 2005
AB 100MG A076900 003 Jan 28, 2005

TENORMIN

AB + ALVOGEN 25MG N018240 004 Apr 09, 1990
AB + 50MG N018240 001
AB +! 100MG N018240 002

ATENOLOL; CHLORTHALIDONE

TABLET;ORAL

ATENOLOL AND CHLORTHALIDONE

AB ALVOGEN 50MG;25MG A072302 002 May 31, 1990
AB 100MG;25MG A072302 001 May 31, 1990
AB MYLAN 50MG;25MG A074203 001 Oct 31, 1993
AB 100MG;25MG A074203 002 Oct 31, 1993
AB WATSON LABS 50MG;25MG A073665 001 Jul 02, 1992
AB 100MG;25MG A073665 002 Jul 02, 1992
AB ZYDUS PHARMS 50MG;25MG A210028 001 Mar 08, 2019
AB 100MG;25MG A210028 002 Mar 08, 2019

TENORETIC 100

AB +! ALVOGEN 100MG;25MG N018760 001 Jun 08, 1984

TENORETIC 50

AB + ALVOGEN 50MG;25MG N018760 002 Jun 08, 1984

PRESCRIPTION DRUG PRODUCT LIST

ATOMOXETINE HYDROCHLORIDE

CAPSULE; ORAL

ATOMOXETINE HYDROCHLORIDE

<u>AB</u>	APOTEX	<u>10MG</u>	<u>A078983 001</u>	May 30, 2017
<u>AB</u>		<u>18MG</u>	<u>A078983 002</u>	May 30, 2017
<u>AB</u>		<u>25MG</u>	<u>A078983 003</u>	May 30, 2017
<u>AB</u>		<u>40MG</u>	<u>A078983 004</u>	May 30, 2017
<u>AB</u>		<u>60MG</u>	<u>A078983 005</u>	May 30, 2017
<u>AB</u>		<u>80MG</u>	<u>A078983 006</u>	May 30, 2017
<u>AB</u>		<u>100MG</u>	<u>A078983 007</u>	May 30, 2017
<u>AB</u>	AUROBINDO PHARMA LTD	<u>10MG</u>	<u>A079016 001</u>	May 30, 2017
<u>AB</u>		<u>18MG</u>	<u>A079016 002</u>	May 30, 2017
<u>AB</u>		<u>25MG</u>	<u>A079016 003</u>	May 30, 2017
<u>AB</u>		<u>40MG</u>	<u>A079016 004</u>	May 30, 2017
<u>AB</u>		<u>60MG</u>	<u>A079016 005</u>	May 30, 2017
<u>AB</u>		<u>80MG</u>	<u>A079016 006</u>	May 30, 2017
<u>AB</u>		<u>100MG</u>	<u>A079016 007</u>	May 30, 2017
<u>AB</u>	DR REDDYS LABS LTD	<u>10MG</u>	<u>A090609 001</u>	Feb 23, 2018
<u>AB</u>		<u>18MG</u>	<u>A090609 002</u>	Feb 23, 2018
<u>AB</u>		<u>25MG</u>	<u>A090609 003</u>	Feb 23, 2018
<u>AB</u>		<u>40MG</u>	<u>A090609 004</u>	Feb 23, 2018
<u>AB</u>		<u>60MG</u>	<u>A090609 005</u>	Feb 23, 2018
<u>AB</u>		<u>80MG</u>	<u>A090609 006</u>	Feb 23, 2018
<u>AB</u>		<u>100MG</u>	<u>A090609 007</u>	Feb 23, 2018
<u>AB</u>	GLENMARK PHARMS LTD	<u>10MG</u>	<u>A079019 001</u>	May 30, 2017
<u>AB</u>		<u>18MG</u>	<u>A079019 002</u>	May 30, 2017
<u>AB</u>		<u>25MG</u>	<u>A079019 003</u>	May 30, 2017
<u>AB</u>		<u>40MG</u>	<u>A079019 004</u>	May 30, 2017
<u>AB</u>		<u>60MG</u>	<u>A079019 005</u>	May 30, 2017
<u>AB</u>		<u>80MG</u>	<u>A079019 006</u>	May 30, 2017
<u>AB</u>		<u>100MG</u>	<u>A079019 007</u>	May 30, 2017
<u>AB</u>	TEVA PHARMS USA	<u>10MG</u>	<u>A079022 001</u>	May 30, 2017
<u>AB</u>		<u>18MG</u>	<u>A079022 002</u>	May 30, 2017
<u>AB</u>		<u>25MG</u>	<u>A079022 003</u>	May 30, 2017
<u>AB</u>		<u>40MG</u>	<u>A079022 004</u>	May 30, 2017
<u>AB</u>		<u>60MG</u>	<u>A079022 005</u>	May 30, 2017
<u>AB</u>		<u>80MG</u>	<u>A079022 006</u>	May 30, 2017
<u>AB</u>		<u>100MG</u>	<u>A079022 007</u>	May 30, 2017
<u>AB</u>	ZYDUS PHARMS USA INC	<u>18MG</u>	<u>A079017 001</u>	Sep 17, 2010
<u>AB</u>		<u>25MG</u>	<u>A079017 002</u>	Sep 17, 2010
<u>AB</u>		<u>40MG</u>	<u>A079017 003</u>	Sep 17, 2010
<u>AB</u>		<u>60MG</u>	<u>A079017 004</u>	Sep 17, 2010
<u>AB</u>		<u>80MG</u>	<u>A079017 005</u>	Sep 17, 2010
<u>AB</u>		<u>100MG</u>	<u>A079017 006</u>	Sep 17, 2010
<u>STRATTERA</u>				
<u>AB</u>	+ LILLY	<u>10MG</u>	<u>N021411 002</u>	Nov 26, 2002
<u>AB</u>	+	<u>18MG</u>	<u>N021411 003</u>	Nov 26, 2002
<u>AB</u>	+	<u>25MG</u>	<u>N021411 004</u>	Nov 26, 2002
<u>AB</u>	+	<u>40MG</u>	<u>N021411 005</u>	Nov 26, 2002
<u>AB</u>	+!	<u>60MG</u>	<u>N021411 006</u>	Nov 26, 2002
<u>AB</u>	+	<u>80MG</u>	<u>N021411 007</u>	Feb 14, 2005
<u>AB</u>	+	<u>100MG</u>	<u>N021411 008</u>	Feb 14, 2005

ATORVASTATIN CALCIUM

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>	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>	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

PRESCRIPTION DRUG PRODUCT LIST

ATORVASTATIN CALCIUM

TABLET;ORAL

ATORVASTATIN CALCIUM

AB		<u>EQ 80MG BASE</u>	<u>A209912 004</u>	Jun 18, 2018
AB	INVAGEN PHARMS	<u>EQ 10MG BASE</u>	<u>A204846 001</u>	Jan 09, 2017
AB		<u>EQ 20MG BASE</u>	<u>A204846 002</u>	Jan 09, 2017
AB		<u>EQ 40MG BASE</u>	<u>A204846 003</u>	Jan 09, 2017
AB		<u>EQ 80MG BASE</u>	<u>A204846 004</u>	Jan 09, 2017
AB	LANNETT CO INC	<u>EQ 10MG BASE</u>	<u>A091624 001</u>	Apr 05, 2013
AB		<u>EQ 20MG BASE</u>	<u>A091624 002</u>	Apr 05, 2013
AB		<u>EQ 40MG BASE</u>	<u>A091624 003</u>	Apr 05, 2013
AB		<u>EQ 80MG BASE</u>	<u>A091624 004</u>	Apr 05, 2013
AB	LUPIN LTD	<u>EQ 10MG BASE</u>	<u>A204991 001</u>	Mar 06, 2019
AB		<u>EQ 20MG BASE</u>	<u>A204991 002</u>	Mar 06, 2019
AB		<u>EQ 40MG BASE</u>	<u>A204991 003</u>	Mar 06, 2019
AB		<u>EQ 80MG BASE</u>	<u>A204991 004</u>	Mar 06, 2019
AB	MICRO LABS LTD INDIA	<u>EQ 10MG BASE</u>	<u>A205945 001</u>	Nov 07, 2019
AB		<u>EQ 20MG BASE</u>	<u>A205945 002</u>	Nov 07, 2019
AB		<u>EQ 40MG BASE</u>	<u>A205945 003</u>	Nov 07, 2019
AB		<u>EQ 80MG BASE</u>	<u>A205945 004</u>	Nov 07, 2019
AB	MSN	<u>EQ 10MG BASE</u>	<u>A211933 001</u>	Feb 08, 2019
AB		<u>EQ 20MG BASE</u>	<u>A211933 002</u>	Feb 08, 2019
AB		<u>EQ 40MG BASE</u>	<u>A211933 003</u>	Feb 08, 2019
AB		<u>EQ 80MG BASE</u>	<u>A211933 004</u>	Feb 08, 2019
AB	MYLAN PHARMS INC	<u>EQ 10MG BASE</u>	<u>A091226 001</u>	May 29, 2012
AB		<u>EQ 20MG BASE</u>	<u>A091226 002</u>	May 29, 2012
AB		<u>EQ 40MG BASE</u>	<u>A091226 003</u>	May 29, 2012
AB		<u>EQ 80MG BASE</u>	<u>A091226 004</u>	May 29, 2012
AB	SANDOZ INC	<u>EQ 10MG BASE</u>	<u>A077575 001</u>	May 29, 2012
AB		<u>EQ 20MG BASE</u>	<u>A077575 002</u>	May 29, 2012
AB		<u>EQ 40MG BASE</u>	<u>A077575 003</u>	May 29, 2012
AB		<u>EQ 80MG BASE</u>	<u>A077575 004</u>	May 29, 2012
AB	SCIEGEN PHARMS INC	<u>EQ 10MG BASE</u>	<u>A205519 001</u>	May 19, 2016
AB		<u>EQ 20MG BASE</u>	<u>A205519 002</u>	May 19, 2016
AB		<u>EQ 40MG BASE</u>	<u>A205519 003</u>	May 19, 2016
AB		<u>EQ 80MG BASE</u>	<u>A205519 004</u>	May 19, 2016
AB	SUN PHARM INDS LTD	<u>EQ 10MG BASE</u>	<u>A076477 001</u>	Nov 30, 2011
AB		<u>EQ 20MG BASE</u>	<u>A076477 002</u>	Nov 30, 2011
AB		<u>EQ 40MG BASE</u>	<u>A076477 003</u>	Nov 30, 2011
AB		<u>EQ 80MG BASE</u>	<u>A076477 004</u>	Nov 30, 2011
AB	TEVA PHARMS USA	<u>EQ 10MG BASE</u>	<u>A205300 001</u>	Mar 27, 2017
AB		<u>EQ 20MG BASE</u>	<u>A205300 002</u>	Mar 27, 2017
AB		<u>EQ 40MG BASE</u>	<u>A205300 003</u>	Mar 27, 2017
AB		<u>EQ 80MG BASE</u>	<u>A205300 004</u>	Mar 27, 2017
AB	THEPHARMANETWORK LLC	<u>EQ 10MG BASE</u>	<u>A209288 001</u>	Dec 21, 2018
AB		<u>EQ 20MG BASE</u>	<u>A209288 002</u>	Dec 21, 2018
AB		<u>EQ 40MG BASE</u>	<u>A209288 003</u>	Dec 21, 2018
AB		<u>EQ 80MG BASE</u>	<u>A209288 004</u>	Dec 21, 2018
AB	ZYDUS PHARMS	<u>EQ 10MG BASE</u>	<u>A206536 001</u>	Nov 20, 2018
AB		<u>EQ 20MG BASE</u>	<u>A206536 002</u>	Nov 20, 2018
AB		<u>EQ 40MG BASE</u>	<u>A206536 003</u>	Nov 20, 2018
AB		<u>EQ 80MG BASE</u>	<u>A206536 004</u>	Nov 20, 2018
<u>LIPITOR</u>				
AB	+ PFIZER	<u>EQ 10MG BASE</u>	<u>N020702 001</u>	Dec 17, 1996
AB	+	<u>EQ 20MG BASE</u>	<u>N020702 002</u>	Dec 17, 1996
AB	+	<u>EQ 40MG BASE</u>	<u>N020702 003</u>	Dec 17, 1996
AB	+	<u>EQ 80MG BASE</u>	<u>N020702 004</u>	Apr 07, 2000

ATORVASTATIN CALCIUM; EZETIMIBE

TABLET;ORAL

EZETIMIBE AND ATORVASTATIN CALCIUM

	WATSON LABS TEVA	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

PRESCRIPTION DRUG PRODUCT LIST

ATOVAQUONE

SUSPENSION; ORAL

ATOVAQUONE

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

MEPRON

AB	+ !	GLAXOSMITHKLINE LLC	<u>750MG/5ML</u>	<u>N020500</u>	<u>001</u>	Feb 08, 1995
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ATOVAQUONE; PROGUANIL HYDROCHLORIDE

TABLET; ORAL

ATOVAQUONE AND PROGUANIL HYDROCHLORIDE

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

MALARONE

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

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

INJECTABLE; INJECTION

ATRAURIUM BESYLATE

AP	AUROBINDO PHARMA LTD	<u>10MG/ML</u>	<u>A206011</u>	<u>001</u>	Apr 08, 2015	
AP	HOSPIRA INC	<u>10MG/ML</u>	<u>A090761</u>	<u>001</u>	Oct 18, 2012	
AP	MEITHEAL	<u>10MG/ML</u>	<u>A091489</u>	<u>001</u>	Feb 17, 2012	
AP	!	WEST-WARD PHARMS INT	<u>10MG/ML</u>	<u>A074901</u>	<u>001</u>	Jul 18, 1997

ATRAURIUM BESYLATE PRESERVATIVE FREE

AP	AUROBINDO PHARMA LTD	<u>10MG/ML</u>	<u>A206010</u>	<u>001</u>	Apr 08, 2015	
AP	HOSPIRA INC	<u>10MG/ML</u>	<u>A090782</u>	<u>001</u>	Oct 18, 2012	
AP	MEITHEAL	<u>10MG/ML</u>	<u>A091488</u>	<u>001</u>	Feb 17, 2012	
AP	!	WEST-WARD PHARMS INT	<u>10MG/ML</u>	<u>A074900</u>	<u>001</u>	Jul 18, 1997

ATROPINE

SOLUTION; INTRAMUSCULAR

ATROPEN

+ !	MERIDIAN MEDCL TECHN	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 SULFATE

SOLUTION; INTRAVENOUS

ATROPINE SULFATE LIFESHIELD ABBOJECT SYRINGE

+ !	HOSPIRA	0.5MG/5ML (0.1MG/ML)	N021146	004	Aug 17, 2017
+ !		1MG/10ML (0.1MG/ML)	N021146	005	Aug 17, 2017

SOLUTION; INTRAVENOUS, INTRAMUSCULAR, SUBCUTANEOUS, ENDOTRACHEAL

ATROPINE SULFATE ANSYR PLASTIC SYRINGE

+ !	HOSPIRA	0.25MG/5ML (0.05MG/ML)	N021146	002	Jul 09, 2001
+ !		1MG/10ML (0.1MG/ML)	N021146	003	Jul 09, 2001

SOLUTION; INTRAVENOUS, INTRAMUSCULAR, SUBCUTANEOUS, INTRAOSSEOUS, ENDOTRACHEAL

ATROPINE SULFATE

+ !	FRESENIUS KABI USA	8MG/20ML (0.4MG/ML)	N209260	001	Jan 26, 2018
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SOLUTION/DROPS; OPHTHALMIC

ATROPINE SULFATE

+ !	AKORN	1%	N206289	001	Jul 18, 2014
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ISOPTO ATROPINE

+ !	ALCON LABS INC	1%	N208151	001	Dec 01, 2016
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ATROPINE SULFATE; DIFENOXIN HYDROCHLORIDE

TABLET; ORAL

MOTOFEN

+ !	SEBELA IRELAND LTD	0.025MG;1MG	N017744	002	
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PRESCRIPTION DRUG PRODUCT LISTATROPINE SULFATE; DIPHENOXYLATE HYDROCHLORIDE

SOLUTION; ORAL

DIPHENOXYLATE HYDROCHLORIDE AND ATROPINE SULFATE
! HIKMA 0.025MG/5ML; 2.5MG/5ML

A087708 001 May 03, 1982

TABLET; ORAL

DIPHENOXYLATE HYDROCHLORIDE AND ATROPINE SULFATE

AA	ANI PHARMS INC	<u>0.025MG; 2.5MG</u>	<u>A086727</u>	<u>001</u>	
AA	BAYSHORE PHARMS LLC	<u>0.025MG; 2.5MG</u>	<u>A210819</u>	<u>001</u>	Nov 13, 2018
AA	LANNETT	<u>0.025MG; 2.5MG</u>	<u>A085372</u>	<u>001</u>	
AA	LEADING PHARMA LLC	<u>0.025MG; 2.5MG</u>	<u>A213413</u>	<u>001</u>	Feb 20, 2020
AA	MYLAN	<u>0.025MG; 2.5MG</u>	<u>A085762</u>	<u>001</u>	
AA	PAR PHARM	<u>0.025MG; 2.5MG</u>	<u>A040357</u>	<u>001</u>	May 02, 2000
AA	UPSHER SMITH LABS	<u>0.025MG; 2.5MG</u>	<u>A210571</u>	<u>001</u>	Aug 31, 2018

LOMOTIL

AA	+! GD SEARLE LLC	<u>0.025MG; 2.5MG</u>	<u>N012462</u>	<u>001</u>	
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ATROPINE; PRALIDOXIME CHLORIDE

INJECTABLE; INTRAMUSCULAR

DUODOTE

+! MERIDIAN MEDCL 2.1MG/0.7ML; 600MG/2ML

N021983 001 Sep 28, 2006

AURANOFIN

CAPSULE; ORAL

RIDAURA

+! SEBELA IRELAND LTD 3MG

N018689 001 May 24, 1985

AVANAFIL

TABLET; ORAL

STENDRA

+ METUCHEN PHARMS 50MG

N202276 001 Apr 27, 2012

+ 100MG

N202276 002 Apr 27, 2012

+! 200MG

N202276 003 Apr 27, 2012

AVAPRITINIB

TABLET; ORAL

AYVAKIT

+ BLUEPRINT MEDICINES 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; IV (INFUSION)

AVYCAZ

+! ALLERGAN 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	<u>100MG/VIAL</u>	<u>A207475</u>	<u>001</u>	Jul 02, 2018
AP	ACTAVIS LLC	<u>100MG/VIAL</u>	<u>N208216</u>	<u>001</u>	Apr 29, 2016
AP	CIPLA	<u>100MG/VIAL</u>	<u>A209540</u>	<u>001</u>	May 04, 2018
AP	DR REDDYS	<u>100MG/VIAL</u>	<u>A201537</u>	<u>001</u>	Sep 16, 2013
AP	MYLAN INSTITUTIONAL	<u>100MG/VIAL</u>	<u>A204949</u>	<u>001</u>	Apr 28, 2016
AP	NATCO PHARMA LTD	<u>100MG/VIAL</u>	<u>A207234</u>	<u>001</u>	Jun 23, 2017
AP	SHILPA MEDICARE	<u>100MG/VIAL</u>	<u>A207518</u>	<u>001</u>	Sep 29, 2016

VIDAZA

AP	+! CELGENE	<u>100MG/VIAL</u>	<u>N050794</u>	<u>001</u>	May 19, 2004
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AZATHIOPRINE

TABLET; ORAL

AZASAN

AB	AAIPHARMA LLC	<u>25MG</u>	<u>A075252</u>	<u>002</u>	Feb 03, 2003
AB		<u>50MG</u>	<u>A075252</u>	<u>001</u>	Jun 07, 1999
AB		<u>75MG</u>	<u>A075252</u>	<u>003</u>	Feb 03, 2003
AB		<u>100MG</u>	<u>A075252</u>	<u>004</u>	Feb 03, 2003

PRESCRIPTION DRUG PRODUCT LIST

3-48 (of 453)

AZATHIOPRINE

TABLET; ORAL

AZATHIOPRINE

<u>AB</u>	AMNEAL PHARMS LLC	<u>50MG</u>	<u>A074069</u>	<u>001</u>	Feb 16, 1996
<u>AB</u>	MYLAN	<u>50MG</u>	<u>A075568</u>	<u>001</u>	Dec 13, 1999
<u>AB</u>	ZYDUS PHARMS USA	<u>25MG</u>	<u>A077621</u>	<u>002</u>	Sep 05, 2008
<u>AB</u>		<u>50MG</u>	<u>A077621</u>	<u>001</u>	Mar 15, 2007
<u>AB</u>		<u>75MG</u>	<u>A077621</u>	<u>003</u>	Sep 05, 2008
<u>AB</u>		<u>100MG</u>	<u>A077621</u>	<u>004</u>	Sep 05, 2008

IMURAN

<u>AB</u>	+! SEBELA IRELAND LTD	<u>50MG</u>	<u>N016324</u>	<u>001</u>	
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AZATHIOPRINE SODIUM

INJECTABLE; INJECTION

AZATHIOPRINE SODIUM

!	WEST-WARD PHARMS INT	EQ 100MG BASE/VIAL	A074419	001	Mar 31, 1995
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AZELAIC ACID

AEROSOL, FOAM; TOPICAL

FINACEA

+	LEO PHARMA AS	15%	N207071	001	Jul 29, 2015
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CREAM; TOPICAL

AZELEX

+	ALMIRALL	20%	N020428	001	Sep 13, 1995
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GEL; TOPICAL

AZELAIC ACID

<u>AB</u>	ACTAVIS LABS UT INC	<u>15%</u>	<u>A208011</u>	<u>001</u>	Nov 19, 2018
<u>AB</u>	GLENMARK PHARMS	<u>15%</u>	<u>A204637</u>	<u>001</u>	Nov 19, 2018
<u>AB</u>	TARO	<u>15%</u>	<u>A210549</u>	<u>001</u>	Aug 23, 2019
<u>AB</u>	TOLMAR	<u>15%</u>	<u>A208724</u>	<u>001</u>	Nov 19, 2018

FINACEA

<u>AB</u>	+! LEO PHARMA AS	<u>15%</u>	<u>N021470</u>	<u>001</u>	Dec 24, 2002
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AZELASTINE HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

AZELASTINE HYDROCHLORIDE

<u>AT</u>	AKORN	<u>0.05%</u>	<u>A203660</u>	<u>001</u>	Nov 08, 2016
<u>AT</u>	ALEMBIC PHARMS LTD	<u>0.05%</u>	<u>A209620</u>	<u>001</u>	Mar 20, 2019
<u>AT</u>	APOTEX INC	<u>0.05%</u>	<u>A078621</u>	<u>001</u>	Aug 03, 2009
<u>AT</u>	GLAND PHARMA LTD	<u>0.05%</u>	<u>A210092</u>	<u>001</u>	Feb 25, 2020
<u>AT</u>	! SANDOZ INC	<u>0.05%</u>	<u>A202305</u>	<u>001</u>	May 31, 2012
<u>AT</u>	SOMERSET THERAPS LLC	<u>0.05%</u>	<u>A207411</u>	<u>001</u>	Mar 29, 2019
<u>AT</u>	SUN PHARM	<u>0.05%</u>	<u>A078738</u>	<u>001</u>	Jun 21, 2010

SPRAY, METERED; NASAL

ASTELIN

<u>AB</u>	+! MYLAN SPECIALITY LP	<u>EQ 0.125MG BASE/SPRAY</u>	<u>N020114</u>	<u>001</u>	Nov 01, 1996
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ASTEPRO

<u>AB</u>	+! MYLAN SPECIALITY LP	<u>EQ 0.1876MG BASE/SPRAY</u>	<u>N022203</u>	<u>002</u>	Aug 31, 2009
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AZELASTINE HYDROCHLORIDE

<u>AB</u>	ALKEM LABS LTD	<u>EQ 0.125MG BASE/SPRAY</u>	<u>A208156</u>	<u>001</u>	Aug 18, 2017
<u>AB</u>	AMNEAL PHARMS LLC	<u>EQ 0.125MG BASE/SPRAY</u>	<u>A204660</u>	<u>001</u>	Aug 28, 2017
<u>AB</u>		<u>EQ 0.1876MG BASE/SPRAY</u>	<u>A208199</u>	<u>001</u>	Dec 15, 2017
<u>AB</u>	APOTEX INC	<u>EQ 0.125MG BASE/SPRAY</u>	<u>A077954</u>	<u>001</u>	Apr 30, 2009
<u>AB</u>		<u>EQ 0.1876MG BASE/SPRAY</u>	<u>A201846</u>	<u>001</u>	Aug 31, 2012
<u>AB</u>	BRECKENRIDGE	<u>EQ 0.125MG BASE/SPRAY</u>	<u>A090176</u>	<u>001</u>	Jul 28, 2015
<u>AB</u>	HI TECH	<u>EQ 0.125MG BASE/SPRAY</u>	<u>A207610</u>	<u>001</u>	May 17, 2019
<u>AB</u>		<u>EQ 0.1876MG BASE/SPRAY</u>	<u>A210032</u>	<u>001</u>	Aug 23, 2019
<u>AB</u>	HIKMA	<u>EQ 0.125MG BASE/SPRAY</u>	<u>A091444</u>	<u>001</u>	Oct 24, 2014
<u>AB</u>		<u>EQ 0.1876MG BASE/SPRAY</u>	<u>A207243</u>	<u>001</u>	Sep 22, 2017
<u>AB</u>	PERRIGO ISRAEL	<u>EQ 0.1876MG BASE/SPRAY</u>	<u>A202743</u>	<u>001</u>	May 08, 2014
<u>AB</u>	SUN PHARM	<u>EQ 0.125MG BASE/SPRAY</u>	<u>A090423</u>	<u>001</u>	May 23, 2012
<u>AB</u>	UPSHER SMITH LABS	<u>EQ 0.125MG BASE/SPRAY</u>	<u>A202609</u>	<u>001</u>	Mar 17, 2017
<u>AB</u>	ZYDUS PHARMS	<u>EQ 0.125MG BASE/SPRAY</u>	<u>A091409</u>	<u>001</u>	Aug 14, 2017

AZELASTINE HYDROCHLORIDE; FLUTICASONE PROPIONATE

SPRAY, METERED; NASAL

AZELASTINE HYDROCHLORIDE AND FLUTICASONE PROPIONATE

<u>AB</u>	APOTEX	<u>EQ 0.125MG BASE/SPRAY; 0.05MG/SPRAY</u>	<u>A207712</u>	<u>001</u>	Apr 28, 2017
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DYMISTA

<u>AB</u>	+! MYLAN SPECIALITY LP	<u>EQ 0.125MG BASE/SPRAY; 0.05MG/SPRAY</u>	<u>N202236</u>	<u>001</u>	May 01, 2012
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PRESCRIPTION DRUG PRODUCT LIST

AZILSARTAN KAMEDOXOMIL

TABLET;ORAL

EDARBI

+	ARBOR PHARMS LLC	EQ 40MG MEDOXOMIL	N200796 001	Feb 25, 2011
+	!	EQ 80MG MEDOXOMIL	N200796 002	Feb 25, 2011

AZILSARTAN KAMEDOXOMIL; CHLORTHALIDONE

TABLET;ORAL

EDARBYCLOR

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

AZITHROMYCIN

FOR SUSPENSION;ORAL

AZITHROMYCIN

<u>AB</u>	AMNEAL PHARMS LLC	<u>EQ 100MG BASE/5ML</u>	<u>A205666 001</u>	Jul 19, 2018
<u>AB</u>		<u>EQ 200MG BASE/5ML</u>	<u>A205666 002</u>	Jul 19, 2018
<u>AB</u>	AUROBINDO PHARMA LTD	<u>EQ 100MG BASE/5ML</u>	<u>A209201 001</u>	Oct 09, 2018
<u>AB</u>		<u>EQ 200MG BASE/5ML</u>	<u>A209201 002</u>	Oct 09, 2018
<u>AB</u>	EPIC PHARMA LLC	<u>EQ 100MG BASE/5ML</u>	<u>A207531 001</u>	Apr 09, 2018
<u>AB</u>		<u>EQ 200MG BASE/5ML</u>	<u>A207531 002</u>	Apr 09, 2018
<u>AB</u>	PLIVA	<u>EQ 100MG BASE/5ML</u>	<u>A065246 002</u>	Jul 05, 2006
<u>AB</u>		<u>EQ 200MG BASE/5ML</u>	<u>A065246 001</u>	Jul 05, 2006
<u>AB</u>	TARO	<u>EQ 200MG BASE/5ML</u>	<u>A211521 001</u>	Dec 11, 2019
<u>AB</u>	TEVA PHARMS	<u>EQ 100MG BASE/5ML</u>	<u>A065419 001</u>	Jun 24, 2008
<u>AB</u>		<u>EQ 200MG BASE/5ML</u>	<u>A065419 002</u>	Jun 24, 2008
<u>AB</u>	ZYDUS	<u>EQ 100MG BASE/5ML</u>	<u>A211147 001</u>	Jul 31, 2018
<u>AB</u>		<u>EQ 200MG BASE/5ML</u>	<u>A211147 002</u>	Jul 31, 2018

ZITHROMAX

<u>AB</u>	+	PFIZER	<u>EQ 100MG BASE/5ML</u>	<u>N050710 001</u>	Oct 19, 1995
<u>AB</u>	+	!	<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

<u>AP</u>	AUROBINDO PHARMA LTD	<u>EQ 500MG BASE/VIAL</u>	<u>A203294 001</u>	Jun 19, 2015
<u>AP</u>	FRESENIUS KABI USA	<u>EQ 500MG BASE/VIAL</u>	<u>A065179 001</u>	Dec 13, 2005
<u>AP</u>	GLAND PHARMA LTD	<u>EQ 500MG BASE/VIAL</u>	<u>A065501 001</u>	Nov 09, 2009
<u>AP</u>	HAINAN POLY PHARM	<u>EQ 500MG BASE/VIAL</u>	<u>A203412 001</u>	Oct 09, 2018
<u>AP</u>	HOSPIRA	<u>EQ 500MG BASE/VIAL</u>	<u>A065500 001</u>	Jun 26, 2009
<u>AP</u>		<u>EQ 500MG BASE/VIAL</u>	<u>A065511 001</u>	Jun 26, 2009
<u>AP</u>	MYLAN ASI	<u>EQ 500MG BASE/VIAL</u>	<u>A065506 001</u>	Mar 24, 2009
<u>AP</u>	MYLAN LABS LTD	<u>EQ 500MG BASE/VIAL</u>	<u>A204732 001</u>	Jan 26, 2017
<u>AP</u>	SUN PHARM INDS LTD	<u>EQ 500MG BASE/VIAL</u>	<u>A090923 001</u>	Apr 02, 2013

ZITHROMAX

<u>AP</u>	+	PFIZER	<u>EQ 500MG BASE/VIAL</u>	<u>N050733 001</u>	Jan 30, 1997
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SOLUTION/DROPS;OPHTHALMIC

AZASITE

+	OAK PHARMS INC	1%	N050810 001	Apr 27, 2007
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TABLET;ORAL

AZITHROMYCIN

<u>AB</u>	ALEMBIC PHARMS LTD	<u>EQ 250MG BASE</u>	<u>A211791 001</u>	Jan 28, 2020
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>A211792 001</u>	Jan 28, 2020
<u>AB</u>		<u>EQ 600MG BASE</u>	<u>A211793 001</u>	Jan 27, 2020
<u>AB</u>	AUROBINDO PHARMA LTD	<u>EQ 250MG BASE</u>	<u>A207370 001</u>	Jul 05, 2018
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>A207398 001</u>	Jul 05, 2018
<u>AB</u>	BIONPHARMA INC	<u>EQ 250MG BASE</u>	<u>A210000 001</u>	Feb 26, 2019
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>A210001 001</u>	Feb 26, 2019
<u>AB</u>		<u>EQ 600MG BASE</u>	<u>A209999 001</u>	Dec 26, 2018
<u>AB</u>	CSPC OUYI	<u>EQ 250MG BASE</u>	<u>A208250 001</u>	Apr 17, 2019
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>A208249 001</u>	Oct 25, 2018
<u>AB</u>		<u>EQ 600MG BASE</u>	<u>A207566 001</u>	Sep 24, 2018
<u>AB</u>	LUPIN LTD	<u>EQ 250MG BASE</u>	<u>A065398 001</u>	May 15, 2015
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>A065399 001</u>	May 15, 2015
<u>AB</u>		<u>EQ 600MG BASE</u>	<u>A065400 001</u>	May 15, 2015
<u>AB</u>	PLIVA	<u>EQ 250MG BASE</u>	<u>A065225 001</u>	Nov 14, 2005
<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>	SUNSHINE LAKE	<u>EQ 250MG BASE</u>	<u>A209045 001</u>	Dec 07, 2018

PRESCRIPTION DRUG PRODUCT LIST

AZITHROMYCIN

TABLET; ORAL

AZITHROMYCIN

AB		<u>EQ 500MG BASE</u>	<u>A209044 001</u>	Dec 07, 2018
AB		<u>EQ 600MG BASE</u>	<u>A209043 001</u>	Dec 06, 2018
AB	TEVA	<u>EQ 500MG BASE</u>	<u>A065193 001</u>	Nov 14, 2005
AB	WOCKHARDT	<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

ZITHROMAX

AB	+	PFIZER	<u>EQ 250MG BASE</u>	<u>N050711 001</u>	Jul 18, 1996
AB	+		<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
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INJECTABLE; INJECTION

AZACTAM

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

AZTREONAM

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

BACITRACIN

INJECTABLE; INJECTION

BACIIM

AP		X GEN PHARMS	<u>50,000 UNITS/VIAL</u>	<u>A064153 001</u>	May 09, 1997
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BACITRACIN

AP		AKORN	<u>50,000 UNITS/VIAL</u>	<u>A206719 001</u>	Oct 20, 2017
AP		FRESENIUS KABI USA	<u>50,000 UNITS/VIAL</u>	<u>A065116 001</u>	Dec 03, 2002
AP	!	PHARMACIA AND UPJOHN	<u>50,000 UNITS/VIAL</u>	<u>A060733 002</u>	
AP		XELLIA PHARMS APS	<u>50,000 UNITS/VIAL</u>	<u>A203177 001</u>	Aug 25, 2014

OINTMENT; OPHTHALMIC

BACITRACIN

!		PERRIGO CO TENNESSEE	500 UNITS/GM	A061212 001	
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BACITRACIN ZINC; HYDROCORTISONE ACETATE; NEOMYCIN SULFATE; POLYMYXIN B SULFATE

OINTMENT; OPHTHALMIC

BACITRACIN-NEOMYCIN-POLYMYXIN W/ HYDROCORTISONE ACETATE

!		PERRIGO CO TENNESSEE	400 UNITS/GM;1%;EQ 3.5MG BASE/GM;10,000 UNITS/GM	A062166 002	
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BACITRACIN ZINC; HYDROCORTISONE; NEOMYCIN SULFATE; POLYMYXIN B SULFATE

OINTMENT; OPHTHALMIC

NEOMYCIN AND POLYMYXIN B SULFATES, BACITRACIN ZINC AND HYDROCORTISONE

AT		AKORN	<u>400 UNITS/GM;1%;EQ 3.5MG BASE/GM;10,000 UNITS/GM</u>	<u>A065213 001</u>	Jul 25, 2012
AT	!	BAUSCH AND LOMB	<u>400 UNITS/GM;1%;EQ 3.5MG BASE/GM;10,000 UNITS/GM</u>	<u>A064068 001</u>	Oct 30, 1995

OINTMENT; TOPICAL

CORTISPORIN

+	!	MONARCH PHARMS	400 UNITS/GM;1%;EQ 3.5MG BASE/GM;5,000 UNITS/GM	N050168 002	May 04, 1984
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BACITRACIN ZINC; NEOMYCIN SULFATE; POLYMYXIN B SULFATE

OINTMENT; OPHTHALMIC

NEOMYCIN AND POLYMYXIN B SULFATES AND BACITRACIN ZINC

AT		AKORN	<u>400 UNITS/GM;EQ 3.5MG BASE/GM;10,000 UNITS/GM</u>	<u>A065088 001</u>	Feb 06, 2004
AT	!	BAUSCH AND LOMB	<u>400 UNITS/GM;EQ 3.5MG BASE/GM;10,000 UNITS/GM</u>	<u>A064064 001</u>	Oct 30, 1995
AT		PERRIGO CO TENNESSEE	<u>400 UNITS/GM;EQ 3.5MG BASE/GM;10,000 UNITS/GM</u>	<u>A060764 002</u>	

NEOSPORIN

AT	+	CASPER PHARMA LLC	<u>400 UNITS/GM;EQ 3.5MG BASE/GM;10,000 UNITS/GM</u>	<u>N050417 001</u>	
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PRESCRIPTION DRUG PRODUCT LIST

BACITRACIN ZINC; POLYMYXIN B SULFATE

OINTMENT;OPHTHALMIC

BACITRACIN ZINC AND POLYMYXIN B SULFATE

<u>AT</u>	AKORN	<u>500 UNITS/GM;10,000 UNITS/GM</u>	<u>A064028 001</u>	Jan 30, 1995
<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>	PERRIGO CO TENNESSEE	<u>500 UNITS/GM;10,000 UNITS/GM</u>	<u>A065022 001</u>	Feb 27, 2002

BACLOFEN

INJECTABLE;INTRATHECAL

BACLOFEN

<u>AP</u>	EMERALD INTL LTD	<u>0.05MG/ML</u>	<u>A091193 001</u>	May 03, 2016
<u>AP</u>		<u>0.5MG/ML</u>	<u>A091193 002</u>	May 03, 2016
<u>AP</u>		<u>2MG/ML</u>	<u>A091193 003</u>	May 03, 2016
<u>AP</u>	MAIA PHARMS INC	<u>0.5MG/ML</u>	<u>A210048 001</u>	Sep 11, 2019
<u>AP</u>		<u>1MG/ML</u>	<u>A210315 001</u>	Jul 30, 2019
<u>AP</u>		<u>2MG/ML</u>	<u>A210048 002</u>	Sep 11, 2019
<u>AP</u>	MYLAN LABS LTD	<u>0.5MG/ML</u>	<u>A209592 001</u>	Mar 21, 2018
<u>AP</u>		<u>1MG/ML</u>	<u>A209594 001</u>	Mar 06, 2018
<u>AP</u>		<u>2MG/ML</u>	<u>A209592 002</u>	Mar 21, 2018

GABLOFEN

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

LIORESAL

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

SOLUTION;ORAL

OZOBAX

+	! METACEL PHARMS LLC	5MG/5ML	N208193 001	Sep 18, 2019
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TABLET;ORAL

BACLOFEN

<u>AB</u>	EYWA PHARMA	<u>10MG</u>	<u>A211555 001</u>	Feb 01, 2019
<u>AB</u>		<u>20MG</u>	<u>A211555 002</u>	Feb 01, 2019
<u>AB</u>	IMPAX LABS	<u>10MG</u>	<u>A077971 001</u>	Oct 26, 2007
<u>AB</u>		<u>20MG</u>	<u>A077971 002</u>	Oct 26, 2007
<u>AB</u>	IVAX SUB TEVA PHARMS	<u>10MG</u>	<u>A072234 001</u>	Jul 21, 1988
<u>AB</u>	!	<u>20MG</u>	<u>A072235 001</u>	Jul 21, 1988
<u>AB</u>	LANNETT CO INC	<u>10MG</u>	<u>A077241 002</u>	Jul 06, 2007
<u>AB</u>		<u>20MG</u>	<u>A077241 001</u>	Dec 20, 2005
<u>AB</u>	MYLAN PHARMS INC	<u>10MG</u>	<u>A090334 001</u>	Feb 18, 2010
<u>AB</u>		<u>20MG</u>	<u>A090334 002</u>	Feb 18, 2010
<u>AB</u>	NORTHSTAR HLTHCARE	<u>10MG</u>	<u>A078401 002</u>	Sep 18, 2009
<u>AB</u>		<u>20MG</u>	<u>A078401 001</u>	Sep 18, 2009
<u>AB</u>	OXFORD PHARMS	<u>10MG</u>	<u>A077088 002</u>	Oct 31, 2007
<u>AB</u>		<u>20MG</u>	<u>A077088 001</u>	Oct 31, 2007
<u>AB</u>	RUBICON	<u>10MG</u>	<u>A209102 002</u>	Nov 28, 2017
<u>AB</u>		<u>20MG</u>	<u>A209102 003</u>	Nov 28, 2017
<u>AB</u>	UPSHER SMITH LABS	<u>10MG</u>	<u>A074584 001</u>	Aug 19, 1996
<u>AB</u>		<u>20MG</u>	<u>A074584 002</u>	Aug 19, 1996
<u>AB</u>	VINTAGE PHARMS	<u>10MG</u>	<u>A077068 002</u>	Aug 30, 2005
<u>AB</u>		<u>20MG</u>	<u>A077068 001</u>	Aug 30, 2005
<u>AB</u>	ZYDUS	<u>10MG</u>	<u>A211659 001</u>	Nov 23, 2018
<u>AB</u>		<u>20MG</u>	<u>A211659 002</u>	Nov 23, 2018
	RUBICON	5MG	A209102 001	Nov 28, 2017

BALOXAVIR MARBOXIL

TABLET;ORAL

XOFLUZA

+	GENENTECH INC	20MG	N210854 001	Oct 24, 2018
+	!	40MG	N210854 002	Oct 24, 2018

BALSALAZIDE DISODIUM

CAPSULE;ORAL

BALSALAZIDE DISODIUM

<u>AB</u>	APOTEX INC	<u>750MG</u>	<u>A077883 001</u>	Dec 28, 2007
<u>AB</u>	HIKMA	<u>750MG</u>	<u>A077806 001</u>	Dec 28, 2007

COLAZAL

<u>AB</u>	! VALEANT PHARMS INTL	<u>750MG</u>	<u>N020610 001</u>	Jul 18, 2000
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PRESCRIPTION DRUG PRODUCT LISTBARICITINIB

TABLET; ORAL

OLUMIANT

+ ELI LILLY AND CO

1MG

N207924 002 Oct 08, 2019

+!

2MG

N207924 001 May 31, 2018

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

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

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

BECLOMETHASONE DIPROPIONATE MONOHYDRATE

SPRAY, METERED; NASAL

BECONASE AQ

+! GLAXOSMITHKLINE

EQ 0.042MG DIPROP/SPRAY

N019389 001 Jul 27, 1987

BEDAQUILINE FUMARATE

TABLET; ORAL

SIRTURO

+! JANSSEN THERAP

EQ 100MG BASE

N204384 001 Dec 28, 2012

BELINOSTAT

POWDER; INTRAVENOUS

BELEODAQ

+! ACROTECH

500MG/VIAL

N206256 001 Jul 03, 2014

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

PRESCRIPTION DRUG PRODUCT LIST

3-53 (of 453)

BENAZEPRIL HYDROCHLORIDE

TABLET; ORAL

BENAZEPRIL HYDROCHLORIDE

<u>AB</u>	AMNEAL PHARMS	<u>5MG</u>	<u>A076820 001</u>	Feb 03, 2006
<u>AB</u>		<u>10MG</u>	<u>A076820 002</u>	Feb 03, 2006
<u>AB</u>		<u>20MG</u>	<u>A076820 003</u>	Feb 03, 2006
<u>AB</u>		<u>40MG</u>	<u>A076820 004</u>	Feb 03, 2006
<u>AB</u>	AUROBINDO PHARMA	<u>10MG</u>	<u>A078212 001</u>	May 22, 2008
<u>AB</u>		<u>20MG</u>	<u>A078212 002</u>	May 22, 2008
<u>AB</u>		<u>40MG</u>	<u>A078212 003</u>	May 22, 2008
<u>AB</u>	CASI PHARMS INC	<u>5MG</u>	<u>A076402 001</u>	Feb 11, 2004
<u>AB</u>		<u>10MG</u>	<u>A076402 002</u>	Feb 11, 2004
<u>AB</u>		<u>20MG</u>	<u>A076402 003</u>	Feb 11, 2004
<u>AB</u>		<u>40MG</u>	<u>A076402 004</u>	Feb 11, 2004
<u>AB</u>	COREPHARMA	<u>5MG</u>	<u>A077128 001</u>	Mar 08, 2006
<u>AB</u>		<u>10MG</u>	<u>A077128 002</u>	Mar 08, 2006
<u>AB</u>		<u>20MG</u>	<u>A077128 003</u>	Mar 08, 2006
<u>AB</u>		<u>40MG</u>	<u>A077128 004</u>	Mar 08, 2006
<u>AB</u>	MYLAN	<u>5MG</u>	<u>A076430 001</u>	Feb 11, 2004
<u>AB</u>		<u>10MG</u>	<u>A076430 002</u>	Feb 11, 2004
<u>AB</u>		<u>20MG</u>	<u>A076430 003</u>	Feb 11, 2004
<u>AB</u>		<u>40MG</u>	<u>A076430 004</u>	Feb 11, 2004
<u>AB</u>	PRINSTON INC	<u>5MG</u>	<u>A076118 001</u>	Feb 11, 2004
<u>AB</u>		<u>10MG</u>	<u>A076118 002</u>	Feb 11, 2004
<u>AB</u>		<u>20MG</u>	<u>A076118 003</u>	Feb 11, 2004
<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 PHARMS USA	<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>	+ US PHARMS HOLDINGS I	<u>5MG</u>	<u>N019851 001</u>	Jun 25, 1991
<u>AB</u>	+	<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>	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>	MYLAN	<u>5MG; 6.25MG</u>	<u>A076688 001</u>	Feb 11, 2004
<u>AB</u>		<u>10MG; 12.5MG</u>	<u>A076688 002</u>	Feb 11, 2004
<u>AB</u>		<u>20MG; 12.5MG</u>	<u>A076688 003</u>	Feb 11, 2004
<u>AB</u>		<u>20MG; 25MG</u>	<u>A076688 004</u>	Feb 11, 2004
<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>	+ US PHARMS HOLDINGS I	<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; IV (INFUSION)

TREANDA

+	CEPHALON	25MG/VIAL	N022249 002	May 01, 2009
+		100MG/VIAL	N022249 001	Mar 20, 2008

PRESCRIPTION DRUG PRODUCT LISTBENDAMUSTINE HYDROCHLORIDE

SOLUTION;IV (INFUSION)

BELRAPZO

+! EAGLE PHARMS 100MG/4ML (25MG/ML) N205580 001 May 15, 2018

BENDEKA

+! EAGLE PHARMS 100MG/4ML (25MG/ML) N208194 001 Dec 07, 2015

BENDROFLUMETHIAZIDE; NADOLOL

TABLET;ORAL

CORZIDE

AB + KING PHARMS LLC **5MG;40MG** **N018647 001** May 25, 1983
AB +! **5MG;80MG** **N018647 002** May 25, 1983

NADOLOL AND BENDROFLUMETHIAZIDE

AB IMPAX LABS **5MG;40MG** **A077833 001** Mar 30, 2007
AB **5MG;80MG** **A077833 002** Mar 30, 2007

BENOXINATE HYDROCHLORIDE; FLUORESCEIN SODIUM

SOLUTION/DROPS;OPHTHALMIC

ALTAFLUOR BENOX

+! ALTAIRE PHARMS INC 0.4%;0.25% N208582 001 Dec 14, 2017

BENZNIDAZOLE

TABLET;ORAL

BENZNIDAZOLE

+ CHEMO RESEARCH SL 12.5MG N209570 001 Aug 29, 2017

+! 100MG N209570 002 Aug 29, 2017

BENZONATATE

CAPSULE;ORAL

BENZONATATE

AA APOTEX INC **100MG** **A091310 001** Jan 16, 2015
AA **200MG** **A091310 002** Jan 16, 2015
AA ASCENT PHARMS INC **100MG** **A211518 001** Feb 22, 2019
AA **150MG** **A211518 002** Feb 22, 2019
AA **200MG** **A211518 003** Feb 22, 2019
AA BIONPHARMA INC **100MG** **A081297 001** Jan 29, 1993
AA **200MG** **A081297 002** Oct 30, 2007
AA CSPC OUYI **100MG** **A202765 002** Aug 25, 2017
AA **200MG** **A202765 001** Jul 31, 2015
AA MIKART **100MG** **A040851 001** Nov 09, 2009
AA **150MG** **A040851 002** Nov 09, 2009
AA **200MG** **A040851 003** Nov 09, 2009
AA ORIT LABS LLC **100MG** **A040682 001** Jul 30, 2007
AA **200MG** **A040682 002** Jul 30, 2007
AA PURACAP PHARM LLC **100MG** **A206948 001** Dec 19, 2018
AA **200MG** **A206948 002** Dec 19, 2018
AA QINGDAO BAHEAL PHARM **100MG** **A210562 001** Nov 09, 2018
AA **150MG** **A210562 002** Nov 09, 2018
AA **200MG** **A210562 003** Nov 09, 2018
AA STRIDES PHARMA **100MG** **A091133 001** Jul 30, 2015
AA **200MG** **A091133 002** Jul 30, 2015
AA ! THEPHARMANETWORK LLC **100MG** **A040627 001** Mar 30, 2007
AA ! **150MG** **A201209 001** Sep 24, 2014
AA ! **200MG** **A040749 001** Jul 25, 2007
AA ZYDUS PHARMS USA **100MG** **A040597 001** Jun 08, 2007
AA **200MG** **A040597 002** Jun 08, 2007

TESSALON**AA** + PFIZER **100MG** **N011210 001**BENZOYL PEROXIDE; CLINDAMYCIN PHOSPHATE

GEL;TOPICAL

ACANYA**AB** +! BAUSCH **2.5%;EQ 1.2% BASE** **N050819 001** Oct 23, 2008BENZACLIN**AB** +! BAUSCH **5%;EQ 1% BASE** **N050756 001** Dec 21, 2000CLINDAMYCIN PHOSPHATE AND BENZOYL PEROXIDE

AB ACTAVIS LABS UT INC **2.5%;EQ 1.2% BASE** **A205128 001** Jun 19, 2015
AB GLENMARK PHARMS **5%;EQ 1% BASE** **A209252 001** Mar 14, 2019
AB MYLAN PHARMS INC **5%;EQ 1% BASE** **A065443 001** Aug 11, 2009
AB PERRIGO ISRAEL **2.5%;EQ 1.2% BASE** **A205397 001** Sep 09, 2019
AB **5%;1.2%** **A090979 001** Jun 26, 2012
AB PERRIGO UK FINCO **5%;EQ 1% BASE** **A202440 001** Sep 21, 2015
AB TARO **2.5%;EQ 1.2% BASE** **A206575 001** Aug 19, 2019

PRESCRIPTION DRUG PRODUCT LIST

3-55 (of 453)

BENZOYL PEROXIDE; CLINDAMYCIN PHOSPHATE

GEL; TOPICAL

CLINDAMYCIN PHOSPHATE AND BENZOYL PEROXIDE

<u>AB</u>		<u>5%;EQ 1% BASE</u>	<u>A208776 001</u>	May 25, 2018
<u>AB</u>	TARO PHARMS	<u>5%;1.2%</u>	<u>A206218 001</u>	Dec 15, 2017
<u>AB</u>	TOLMAR	<u>5%;EQ 1% BASE</u>	<u>A204087 001</u>	Jun 27, 2017
<u>AB</u>		<u>5%;1.2%</u>	<u>A203688 001</u>	Aug 25, 2016
<u>AB</u>	ZYDUS PHARMS	<u>5%;1.2%</u>	<u>A210794 001</u>	Dec 28, 2018
<u>DUAC</u>				
<u>AB</u>	+! STIEFEL	<u>5%;1.2%</u>	<u>N050741 001</u>	Aug 26, 2002
ONEXTON				
	+! BAUSCH	3.75%;EQ 1.2% BASE	N050819 002	Nov 24, 2014

BENZOYL PEROXIDE; ERYTHROMYCIN

GEL; TOPICAL

BENZAMYCIN

<u>AB</u>	+! VALEANT INTL	<u>5%;3%</u>	<u>N050557 001</u>	Oct 26, 1984
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ERYTHROMYCIN AND BENZOYL PEROXIDE

<u>AB</u>	LYNE	<u>5%;3%</u>	<u>A065385 001</u>	Sep 18, 2015
<u>AB</u>	TOLMAR	<u>5%;3%</u>	<u>A065112 001</u>	Mar 29, 2004

BENZPHETAMINE HYDROCHLORIDE

TABLET; ORAL

BENZPHETAMINE HYDROCHLORIDE

<u>AA</u>	ANDA REPOSITORY	<u>50MG</u>	<u>A090473 002</u>	Sep 15, 2010
<u>AA</u>	EMCURE PHARMS LTD	<u>50MG</u>	<u>A202061 001</u>	Jan 27, 2012
<u>AA</u>	EPIC PHARMA LLC	<u>50MG</u>	<u>A090346 001</u>	Dec 15, 2015
<u>AA</u>	! KVK TECH	<u>50MG</u>	<u>A090968 001</u>	Jul 20, 2010
<u>AA</u>	SPECGX LLC	<u>50MG</u>	<u>A040773 001</u>	Apr 25, 2007
<u>AA</u>	TWI PHARMS	<u>50MG</u>	<u>A040578 001</u>	Apr 17, 2006
	ANDA REPOSITORY	25MG	A090473 001	Sep 15, 2010

BENZTROPINE MESYLATE

INJECTABLE; INJECTION

BENZTROPINE MESYLATE

<u>AP</u>	FRESENIUS KABI USA	<u>1MG/ML</u>	<u>A090233 001</u>	Jul 28, 2009
<u>AP</u>	HIKMA FARMACEUTICA	<u>1MG/ML</u>	<u>A090287 001</u>	Aug 31, 2009
<u>AP</u>	NAVINTA LLC	<u>1MG/ML</u>	<u>A091525 001</u>	Feb 05, 2013

COGENTIN

<u>AP</u>	+! OAK PHARMS AKORN	<u>1MG/ML</u>	<u>N012015 001</u>	
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TABLET; ORAL

BENZTROPINE MESYLATE

<u>AA</u>	! ASPEN GLOBAL INC	<u>0.5MG</u>	<u>A204713 001</u>	Apr 14, 2015
<u>AA</u>	!	<u>1MG</u>	<u>A204713 002</u>	Apr 14, 2015
<u>AA</u>	!	<u>2MG</u>	<u>A204713 003</u>	Apr 14, 2015
<u>AA</u>	EPIC PHARMA LLC	<u>0.5MG</u>	<u>A072264 001</u>	Feb 27, 1989
<u>AA</u>		<u>1MG</u>	<u>A072265 001</u>	Feb 27, 1989
<u>AA</u>		<u>2MG</u>	<u>A072266 001</u>	Feb 27, 1989
<u>AA</u>	INVAGEN PHARMS	<u>0.5MG</u>	<u>A090294 001</u>	Mar 29, 2010
<u>AA</u>		<u>1MG</u>	<u>A090294 002</u>	Mar 29, 2010
<u>AA</u>		<u>2MG</u>	<u>A090294 003</u>	Mar 29, 2010
<u>AA</u>	LEADING PHARMA LLC	<u>0.5MG</u>	<u>A090168 001</u>	Nov 28, 2012
<u>AA</u>		<u>1MG</u>	<u>A090168 002</u>	Nov 28, 2012
<u>AA</u>		<u>2MG</u>	<u>A090168 003</u>	Nov 28, 2012
<u>AA</u>	PLIVA	<u>0.5MG</u>	<u>A089058 001</u>	Aug 10, 1988
<u>AA</u>		<u>1MG</u>	<u>A089059 001</u>	Aug 10, 1988
<u>AA</u>		<u>2MG</u>	<u>A089060 001</u>	Aug 10, 1988
<u>AA</u>	VINTAGE	<u>0.5MG</u>	<u>A040715 001</u>	Aug 27, 2007
<u>AA</u>		<u>1MG</u>	<u>A040715 002</u>	Aug 27, 2007
<u>AA</u>		<u>2MG</u>	<u>A040715 003</u>	Aug 27, 2007

BENZYL PENICILLOYL POLYLYSINE

INJECTABLE; INJECTION

PRE-PEN

	+! ALLERQUEST	60UMOLAR	N050114 001	
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BEPOTASTINE BESILATE

SOLUTION/DROPS; OPHTHALMIC

BEPREVE

	+! BAUSCH AND LOMB INC	1.5%	N022288 001	Sep 08, 2009
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PRESCRIPTION DRUG PRODUCT LIST

BERACTANT

SUSPENSION; INTRATRACHEAL

SURVANTA

+! ABBVIE 25MG/ML N020032 001 Jul 01, 1991

BESIFLOXACIN HYDROCHLORIDE

SUSPENSION/DROPS; OPHTHALMIC

BESIVANCE

+! BAUSCH AND LOMB EQ 0.6% BASE N022308 001 May 28, 2009

BETAINE

FOR SOLUTION; ORAL

CYSTADANE

+! ORPHAN EUROPE 1GM/SCOOPFUL N020576 001 Oct 25, 1996

BETAMETHASONE ACETATE; BETAMETHASONE SODIUM PHOSPHATE

INJECTABLE; INJECTION

BETAMETHASONE ACETATE AND BETAMETHASONE SODIUM PHOSPHATEAB AM REGENT 3MG/ML; EQ 3MG BASE/ML A090747 001 Jul 31, 2009CELESTONE SOLUSPANAB +! MERCK SHARP DOHME 3MG/ML; EQ 3MG BASE/ML N014602 001BETAMETHASONE DIPROPIONATE

CREAM; TOPICAL

BETAMETHASONE DIPROPIONATEAB ACP NIMBLE EQ 0.05% BASE A210217 001 Oct 12, 2018AB ACTAVIS MID EQ 0.05% BASE A070885 001 Feb 03, 1987

ATLANTIC

AB +! FOUGERA PHARMS EQ 0.05% BASE N019137 001 Jun 26, 1984AB TARO EQ 0.05% BASE A073552 001 Apr 30, 1992AB ZYDUS PHARMS EQ 0.05% BASE A208885 001 Jan 11, 2019

CREAM, AUGMENTED; TOPICAL

BETAMETHASONE DIPROPIONATEAB ANDA REPOSITORY EQ 0.05% BASE A076603 001 Jan 23, 2004AB FOUGERA PHARMS EQ 0.05% BASE A076215 001 Dec 09, 2003AB ! GLENMARK GENERICS EQ 0.05% BASE A078930 001 Sep 23, 2008AB PERRIGO ISRAEL EQ 0.05% BASE A076592 001 Dec 09, 2003AB TARO EQ 0.05% BASE A076543 001 Dec 09, 2003

GEL, AUGMENTED; TOPICAL

BETAMETHASONE DIPROPIONATEAB ! FOUGERA PHARMS EQ 0.05% BASE A075276 001 May 13, 2003AB TARO EQ 0.05% BASE A076508 001 Dec 02, 2003

LOTION; TOPICAL

BETAMETHASONE DIPROPIONATEAB ACP NIMBLE EQ 0.05% BASE A071467 001 Aug 10, 1987AB ACTAVIS MID EQ 0.05% BASE A070281 001 Jul 31, 1985

ATLANTIC

AB ! FOUGERA PHARMS INC EQ 0.05% BASE A070275 001 Aug 12, 1985AB HI TECH EQ 0.05% BASE A209896 001 Feb 06, 2018AB PERRIGO NEW YORK EQ 0.05% BASE A072538 001 Jan 31, 1990

LOTION, AUGMENTED; TOPICAL

BETAMETHASONE DIPROPIONATEAB FOUGERA PHARMS EQ 0.05% BASE A077111 001 May 21, 2007AB HI TECH EQ 0.05% BASE A208849 001 Oct 11, 2019AB ! TARO EQ 0.05% BASE A077477 001 May 21, 2007AB TELIGENT PHARMA INC EQ 0.05% BASE A206389 001 Feb 13, 2018

OINTMENT; TOPICAL

BETAMETHASONE DIPROPIONATEAB ACTAVIS MID EQ 0.05% BASE A071012 001 Feb 03, 1987

ATLANTIC

AB +! FOUGERA PHARMS INC EQ 0.05% BASE N019141 001 Sep 04, 1984AB TARO EQ 0.05% BASE A074271 001 Sep 15, 1994

OINTMENT, AUGMENTED; TOPICAL

BETAMETHASONE DIPROPIONATEAB ACTAVIS MID EQ 0.05% BASE A074304 001 Aug 31, 1995

ATLANTIC

AB FOUGERA PHARMS EQ 0.05% BASE A075373 001 Jun 22, 1999AB LUPIN LTD EQ 0.05% BASE A209106 001 Dec 18, 2019AB TARO EQ 0.05% BASE A076753 001 Oct 12, 2004AB TELIGENT PHARMA INC EQ 0.05% BASE A206118 001 Nov 09, 2017DIPROLENEAB +! MERCK SHARP DOHME EQ 0.05% BASE N018741 001 Jul 27, 1983

SPRAY; TOPICAL

SERNIVO

+! ENCORE DERMAT 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

OINTMENT;TOPICAL

CALCIPOTRIENE AND BETAMETHASONE DIPROPIONATE**AB** PERRIGO UK FINCO **0.064%;0.005%****A200174 001** Dec 12, 2014**AB** TOLMAR **0.064%;0.005%****A201615 001** Jan 14, 2013TACLONEX**AB** +! LEO PHARMA AS **0.064%;0.005%****N021852 001** Jan 09, 2006

SUSPENSION;TOPICAL

TACLONEX

+! LEO PHARMA AS 0.064%;0.005%

N022185 001 May 09, 2008

BETAMETHASONE 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 PHARMS **EQ 0.05% BASE;1%****A202894 001** Oct 30, 2015**AB** TARO **EQ 0.05% BASE;1%****A075673 001** May 29, 2001LOTRISONE**AB** +! MERCK SHARP DOHME **EQ 0.05% BASE;1%****N018827 001** Jul 10, 1984

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, 2004LOTRISONE**AB** +! MERCK SHARP DOHME **EQ 0.05% BASE;1%****N020010 001** Dec 08, 2000BETAMETHASONE VALERATE

AEROSOL, FOAM;TOPICAL

BETAMETHASONE VALERATE**AB** PERRIGO UK FINCO **0.12%****A078337 001** Nov 26, 2012**AB** RICONPHARMA LLC **0.12%****A207144 001** May 24, 2017**AB** TARO **0.12%****A208204 001** May 24, 2017LUXIQ**AB** +! MYLAN **0.12%****N020934 001** Feb 28, 1999

CREAM;TOPICAL

BETA-VAL**AB** ACP NIMBLE **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** ANIMA **EQ 0.1% BASE****A070052 001** Jul 31, 1985**AB** +! FOUGERA PHARMS INC **EQ 0.1% BASE****N018866 001** Aug 31, 1983

OINTMENT;TOPICAL

BETA-VAL**AB** ACP NIMBLE **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** AKORN **EQ 0.5% BASE****A075386 001** Jun 30, 2000**AT** MEDIMETRIKS PHARMS **EQ 0.5% BASE****A075630 001** Apr 12, 2001**AT** WOCKHARDT **EQ 0.5% BASE****A078694 001** Nov 16, 2009BETOPTIC**AT** +! SANDOZ INC **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>		ECI PHARMS LLC	<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>		HERITAGE PHARMA	<u>5MG</u>	<u>A091256</u>	<u>001</u>	May 04, 2010
<u>AA</u>			<u>10MG</u>	<u>A091256</u>	<u>002</u>	May 04, 2010
<u>AA</u>			<u>25MG</u>	<u>A091256</u>	<u>003</u>	May 04, 2010
<u>AA</u>			<u>50MG</u>	<u>A091256</u>	<u>004</u>	May 04, 2010
<u>AA</u>		LANNETT CO INC	<u>5MG</u>	<u>A040677</u>	<u>002</u>	Mar 27, 2008
<u>AA</u>			<u>10MG</u>	<u>A040677</u>	<u>003</u>	Mar 27, 2008
<u>AA</u>			<u>25MG</u>	<u>A040677</u>	<u>004</u>	Mar 27, 2008
<u>AA</u>			<u>50MG</u>	<u>A040677</u>	<u>001</u>	Mar 27, 2008
<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
<u>AA</u>		WOCKHARDT	<u>5MG</u>	<u>A040532</u>	<u>001</u>	Sep 29, 2003
<u>AA</u>			<u>10MG</u>	<u>A040533</u>	<u>001</u>	Sep 29, 2003
<u>AA</u>			<u>25MG</u>	<u>A040534</u>	<u>001</u>	Sep 29, 2003
<u>AA</u>			<u>50MG</u>	<u>A040518</u>	<u>001</u>	Sep 29, 2003

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>	

URECHOLINE

<u>AA</u>	!	ODYSSEY PHARMS	<u>50MG</u>	<u>A089096</u>	<u>001</u>	Dec 19, 1985
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BETRIXABAN

CAPSULE;ORAL

BEVYXXA

	+	PORTOLA PHARMS INC	40MG	N208383	001	Jun 23, 2017
	+	!	80MG	N208383	002	Jun 23, 2017

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 INC	<u>75MG</u>	<u>A209861</u>	<u>001</u>	May 08, 2018
<u>AB</u>		BIONPHARMA INC	<u>75MG</u>	<u>A203174</u>	<u>001</u>	Aug 12, 2014
<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
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GEL;TOPICAL

	+	!	BAUSCH	1%	N021056	001	Jun 28, 2000
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BICALUTAMIDE

TABLET;ORAL

BICALUTAMIDE

<u>AB</u>		ACCORD HLTHCARE	<u>50MG</u>	<u>A078917</u>	<u>001</u>	Jul 06, 2009
<u>AB</u>		APOTEX INC	<u>50MG</u>	<u>A200274</u>	<u>001</u>	May 21, 2015
<u>AB</u>		KENTON	<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
<u>AB</u>		ZYDUS PHARMS USA INC	<u>50MG</u>	<u>A079089</u>	<u>001</u>	Jul 06, 2009

CASODEX

<u>AB</u>	+	!	ANI PHARMS INC	<u>50MG</u>	<u>N020498</u>	<u>001</u>	Oct 04, 1995
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PRESCRIPTION DRUG PRODUCT LIST

3-59 (of 453)

BICTEGRAVIR SODIUM; EMTRICITABINE; TENOFOVIR ALAFENAMIDE FUMARATE

TABLET;ORAL

BIKTARVY

+! GILEAD SCIENCES INC EQ 50MG BASE;200MG;EQ 25MG BASE N210251 001 Feb 07, 2018

BIMATOPROST

SOLUTION/DROPS;OPHTHALMIC

BIMATOPROST

AT	ALEMBIC PHARMS LTD	0.03%	A210263 001	Apr 12, 2019
AT	APOTEX INC	0.03%	A090449 001	Jul 20, 2015
AT	GLAND PHARMA LTD	0.03%	A210126 001	Mar 22, 2019
AT	! LUPIN LTD	0.03%	A203991 001	Feb 20, 2015
AT	SANDOZ INC	0.03%	A202565 001	May 05, 2015
AT	SOMERSET THERAPS LLC	0.03%	A207601 001	Jun 19, 2019

LUMIGAN

+! ALLERGAN 0.01% N022184 001 Aug 31, 2010

SOLUTION/DROPS;TOPICAL

BIMATOPROST

AT	ALEMBIC PHARMS LTD	0.03%	A210515 001	Jan 21, 2020
AT	APOTEX INC	0.03%	A201894 001	Dec 01, 2014
AT	HI TECH	0.03%	A203051 001	Oct 09, 2018
AT	SANDOZ INC	0.03%	A202719 001	Apr 19, 2016

LATISSE**AT** +! ALLERGAN **0.03%** **N022369 001** Dec 24, 2008BINIMETINIB

TABLET;ORAL

MEKTOVI

+! ARRAY BIOPHARMA INC 15MG N210498 001 Jun 27, 2018

BISMUTH SUBCITRATE POTASSIUM; METRONIDAZOLE; TETRACYCLINE HYDROCHLORIDE

CAPSULE;ORAL

PYLERA

+! ALLERGAN 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	AUROBINDO PHARMA	5MG	A077910 001	Dec 27, 2006
AB		10MG	A077910 002	Dec 27, 2006
AB	CASI PHARMS INC	5MG	A075643 001	Nov 16, 2000
AB		10MG	A075643 002	Nov 16, 2000
AB	FRONTIDA BIOPHARM	5MG	A075474 001	Oct 25, 2002
AB		10MG	A075474 002	Oct 25, 2002
AB	ORIT LABS LLC	5MG	A204891 001	Jan 11, 2017
AB		10MG	A204891 002	Jan 11, 2017
AB	TEVA PHARMS	5MG	A075644 001	Jun 26, 2001
AB		10MG	A075644 002	Jun 26, 2001
AB	UNICHEM PHARMS (USA)	5MG	A078635 001	Aug 18, 2009
AB	!	10MG	A078635 002	Aug 18, 2009

BISOPROLOL FUMARATE; HYDROCHLOROTHIAZIDE

TABLET;ORAL

BISOPROLOL FUMARATE AND HYDROCHLOROTHIAZIDE

AB	SANDOZ	2.5MG;6.25MG	A075579 001	Sep 25, 2000
AB		5MG;6.25MG	A075579 002	Sep 25, 2000
AB		10MG;6.25MG	A075579 003	Sep 25, 2000
AB	UNICHEM	2.5MG;6.25MG	A079106 001	Jul 28, 2010
AB		5MG;6.25MG	A079106 002	Jul 28, 2010
AB		10MG;6.25MG	A079106 003	Jul 28, 2010

ZIAC

AB	+	TEVA BRANDED PHARM	2.5MG;6.25MG	N020186 003	Mar 26, 1993
AB	+		5MG;6.25MG	N020186 001	Mar 26, 1993
AB	+		10MG;6.25MG	N020186 002	Mar 26, 1993

PRESCRIPTION DRUG PRODUCT LIST

BIVALIRUDIN

INJECTABLE; INTRAVENOUS

ANGIOMAX

AP +! SANDOZ INC **250MG/VIAL** **N020873 001** Dec 15, 2000

BIVALIRUDIN

AP ACCORD HLTHCARE **250MG/VIAL** **A206551 001** Nov 22, 2017

AP AUROBINDO PHARMA LTD **250MG/VIAL** **A205962 001** Jul 27, 2018

AP CIPLA **250MG/VIAL** **A091602 001** Jul 16, 2018

AP DR REDDYS LABS LTD **250MG/VIAL** **A201577 001** May 26, 2017

AP FRESENIUS KABI USA **250MG/VIAL** **A090189 001** Oct 28, 2016

AP HOSPIRA INC **250MG/VIAL** **A090811 001** Jul 14, 2015

AP **250MG/VIAL** **A090816 001** Jul 14, 2015

AP MYLAN INSTITUTIONAL **250MG/VIAL** **A202471 001** Jun 01, 2018

AP SHUANGCHENG **250MG/VIAL** **A210031 001** Oct 23, 2019

SOLUTION; INTRAVENOUS

ANGIOMAX RTU

+! MAIA PHARMS INC 250MG/50ML (5MG/ML) N211215 001 Jul 25, 2019

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

BLEOMYCIN SULFATE

AP CIPLA **EQ 15 UNITS BASE/VIAL** **A209439 001** Mar 11, 2019

AP ! FRESENIUS KABI USA **EQ 15 UNITS BASE/VIAL** **A065185 001** Jan 28, 2008

AP ! **EQ 30 UNITS BASE/VIAL** **A065185 002** Jan 28, 2008

AP HOSPIRA **EQ 15 UNITS BASE/VIAL** **A065031 001** Mar 10, 2000

AP **EQ 30 UNITS BASE/VIAL** **A065031 002** Mar 10, 2000

AP MEITHEAL **EQ 15 UNITS BASE/VIAL** **A205030 001** Apr 20, 2018

AP **EQ 30 UNITS BASE/VIAL** **A205030 002** Apr 20, 2018

AP TEVA PHARMS USA **EQ 15 UNITS BASE/VIAL** **A065033 001** Jun 27, 2000

AP **EQ 30 UNITS BASE/VIAL** **A065033 002** Jun 27, 2000

AP WEST-WARD PHARMS INT **EQ 15 UNITS BASE/VIAL** **A065042 002** Oct 17, 2001

AP **EQ 30 UNITS BASE/VIAL** **A065042 001** Oct 17, 2001

BORTEZOMIB

INJECTABLE; INTRAVENOUS, SUBCUTANEOUS

VELCADE

+! MILLENNIUM PHARMS 3.5MG/VIAL N021602 001 May 13, 2003

POWDER; INTRAVENOUS

BORTEZOMIB

DR REDDYS LABS LTD 3.5MG/VIAL N206927 001 Oct 04, 2019

FRESENIUS KABI USA 3.5MG/VIAL N205004 001 Nov 06, 2017

BOSENTAN

TABLET; ORAL

BOSENTAN

AB ALEMbic PHARMS LTD **62.5MG** **A211461 001** Jan 23, 2020

AB **125MG** **A211461 002** Jan 23, 2020

AB AMNEAL PHARMS CO **62.5MG** **A209742 001** Apr 26, 2019

AB **125MG** **A209742 002** Apr 26, 2019

AB CIPLA **62.5MG** **A210342 001** Jan 03, 2020

AB **125MG** **A210342 002** Jan 03, 2020

AB HIKMA **62.5MG** **A208695 001** Apr 26, 2019

AB **125MG** **A208695 002** Apr 26, 2019

AB NATCO PHARMA LTD **62.5MG** **A206987 001** Apr 26, 2019

AB **125MG** **A206987 002** Apr 26, 2019

AB PAR PHARM INC **62.5MG** **A205699 001** Apr 26, 2019

AB **125MG** **A205699 002** Apr 26, 2019

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 PHARMS LTD **62.5MG** **N021290 001** Nov 20, 2001

AB +! **125MG** **N021290 002** Nov 20, 2001

PRESCRIPTION DRUG PRODUCT LIST

BOSENTAN

TABLET, FOR SUSPENSION;ORAL

TRACLEER

+! ACTELION PHARMS 32MG N209279 001 Sep 05, 2017

BOSUTINIB MONOHYDRATE

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)

+! AMAG PHARMS INC EQ 1.75MG BASE/0.3ML (EQ 1.75MG
BASE/0.3 ML) N210557 001 Jun 21, 2019BRETYLIUM TOSYLATE

INJECTABLE;INJECTION

BRETYLIUM TOSYLATE

! BRECKENRIDGE 50MG/ML A204386 001 Dec 21, 2018

BREXANOLONE

SOLUTION;INTRAVENOUS

ZULRESSO

+! SAGE THERAP 100MG/20ML (5MG/ML) N211371 001 Jun 17, 2019

BREXPIPIRAZOLE

TABLET;ORAL

REXULTI

+ OTSUKA PHARM CO LTD 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

+ ARIAD 30MG N208772 001 Apr 28, 2017

+ 90MG N208772 002 Apr 28, 2017

+! 180MG N208772 003 Oct 02, 2017

BRILLIANT BLUE G

SOLUTION;OPHTHALMIC

TISSUEBLUE

+! DUTCH OPHTHALMIC 0.025% N209569 001 Dec 20, 2019

BRIMONIDINE TARTRATE

GEL;TOPICAL

MIRVASO

+! GALDERMA LABS LP EQ 0.33% BASE N204708 001 Aug 23, 2013

SOLUTION/DROPS;OPHTHALMIC

ALPHAGAN P**AT** +! ALLERGAN **0.15%** **N021262 001** Mar 16, 2001BRIMONIDINE TARTRATE**AT** AKORN **0.2%** **A076439 001** Mar 14, 2006**AT** ! BAUSCH AND LOMB **0.2%** **A076260 001** May 28, 2003**AT** INDOCO REMEDIES **0.2%** **A091691 001** Nov 18, 2014**AT** SANDOZ INC **0.2%** **A076254 001** Sep 16, 2003**AT** **0.2%** **A078075 001** Jan 30, 2008**AT** SOMERSET THERAPS **0.2%** **A208992 001** Mar 11, 2019

LLC

QOLIANA**AT** SANDOZ INC **0.15%** **N021764 001** May 22, 2006

ALPHAGAN P

+! ALLERGAN 0.1% N021770 001 Aug 19, 2005

BRIMONIDINE TARTRATE; BRINZOLAMIDE

SUSPENSION/DROPS;OPHTHALMIC

SIMBRINZA

+! NOVARTIS 0.2%;1% N204251 001 Apr 19, 2013

PRESCRIPTION DRUG PRODUCT LIST

BRIMONIDINE TARTRATE; TIMOLOL MALEATE

SOLUTION/DROPS;OPHTHALMIC

COMBIGAN

+! ALLERGAN 0.2%;EQ 0.5% BASE N021398 001 Oct 30, 2007

BRINZOLAMIDE

SUSPENSION/DROPS;OPHTHALMIC

AZOPT

+! NOVARTIS 1% N020816 001 Apr 01, 1998

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 SODIUM**AT2** ALEMBIC PHARMS LTD **EQ 0.09% ACID** **A210560 001** Jun 21, 2019**AT2** GLAND PHARMA LTD **EQ 0.09% ACID** **A211029 001** Mar 17, 2020**AT2** ! HI TECH **EQ 0.09% ACID** **A203395 001** Jan 22, 2014**AT2** MYLAN **EQ 0.09% ACID** **A203368 001** Jun 03, 2019

BROMSITE

+! SUN PHARMA GLOBAL EQ 0.075% ACID N206911 001 Apr 08, 2016

PROLENSA

+! BAUSCH AND LOMB EQ 0.07% ACID N203168 001 Apr 05, 2013

BROMOCRIPTINE MESYLATE

CAPSULE;ORAL

BROMOCRIPTINE MESYLATE**AB** ! MYLAN **EQ 5MG BASE** **A077226 001** Apr 04, 2005**AB** ZYDUS PHARMS USA **EQ 5MG BASE** **A078899 001** Jul 30, 2008

INC

PARLODEL**AB** + US PHARMS HOLDINGS **EQ 5MG BASE** **N017962 002** Mar 01, 1982

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TABLET;ORAL

BROMOCRIPTINE MESYLATE**AB** MYLAN **EQ 2.5MG BASE** **A076962 001** Sep 24, 2004**AB** ! PADDOCK LLC **EQ 2.5MG BASE** **A077646 001** Oct 01, 2008**AB** SANDOZ INC **EQ 2.5MG BASE** **A074631 001** Jan 13, 1998PARLODEL**AB** + US PHARMS HOLDINGS **EQ 2.5MG BASE** **N017962 001**

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CYCLOSET

+! VEROSCIENCE EQ 0.8MG BASE N020866 001 May 05, 2009

BROMPHENIRAMINE MALEATE; DEXTROMETHORPHAN HYDROBROMIDE; PSEUDOEPHEDRINE HYDROCHLORIDE

SYRUP;ORAL

BROMFED-DM**AA** ! WOCKHARDT BIO AG **2MG/5ML;10MG/5ML;30MG/5ML** **A088811 001** Jun 07, 1985BROMPHENIRAMINE MALEATE, PSEUDOEPHEDRINE HYDROCHLORIDE AND DEXTROMETHORPHAN HYDROBROMIDE**AA** ACELLA **2MG/5ML;10MG/5ML;30MG/5ML** **A203375 001** Sep 20, 2016**AA** MAYNE PHARMA INC **2MG/5ML;10MG/5ML;30MG/5ML** **A207676 001** Dec 04, 2018**AA** PADDOCK LLC **2MG/5ML;10MG/5ML;30MG/5ML** **A205292 001** Jul 15, 2014**AA** TARO **2MG/5ML;10MG/5ML;30MG/5ML** **A205112 001** Feb 27, 2017BUDESONIDE

AEROSOL, FOAM;RECTAL

UCERIS

+! SALIX 2MG/ACTUATION N205613 001 Oct 07, 2014

CAPSULE;ORAL

BUDESONIDE**AB** ALVOGEN **3MG** **A206724 001** Nov 23, 2016**AB** AMNEAL PHARMS **3MG** **A206200 001** Jul 31, 2017**AB** APPCO **3MG** **A207367 001** Apr 07, 2017

PRESCRIPTION DRUG PRODUCT LIST

BUDESONIDE

CAPSULE; ORAL

BUDESONIDE

<u>AB</u>	BARR LABS DIV TEVA	<u>3MG</u>	<u>A090379</u>	<u>001</u>	Apr 02, 2014
<u>AB</u>	MAYNE PHARMA	<u>3MG</u>	<u>A206623</u>	<u>001</u>	Apr 08, 2016
<u>AB</u>	MYLAN	<u>3MG</u>	<u>A090410</u>	<u>001</u>	May 16, 2011
<u>AB</u>	SCIECURE PHARMA INC	<u>3MG</u>	<u>A209041</u>	<u>001</u>	Sep 28, 2017
<u>AB</u>	ZYDUS PHARMS	<u>3MG</u>	<u>A206134</u>	<u>001</u>	May 04, 2017

ENTOCORT EC

<u>AB</u>	+!	PERRIGO PHARMA INTL	<u>3MG</u>	<u>N021324</u>	<u>001</u>	Oct 02, 2001
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CAPSULE, EXTENDED RELEASE; ORAL

ORTIKOS

	+	SUN PHARMA GLOBAL	6MG	N211929	001	Jun 13, 2019
	+	!	9MG	N211929	002	Jun 13, 2019

POWDER, METERED; INHALATION

PULMICORT FLEXHALER

	+	ASTRAZENECA	0.08MG/INH	N021949	001	Jul 12, 2006
	+	!	0.16MG/INH	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 ATLANTIS	<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 INC	<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>		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 PHARMS	<u>0.25MG/2ML</u>	<u>N020929</u>	<u>001</u>	Aug 08, 2000
<u>AN</u>	+		<u>0.5MG/2ML</u>	<u>N020929</u>	<u>002</u>	Aug 08, 2000
<u>AN</u>	+	!	<u>1MG/2ML</u>	<u>N020929</u>	<u>003</u>	Aug 08, 2000

TABLET, EXTENDED RELEASE; ORAL

BUDESONIDE

<u>AB</u>		ACTAVIS LABS FL INC	<u>9MG</u>	<u>A205457</u>	<u>001</u>	Jul 03, 2018
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UCERIS

<u>AB</u>	+	!	SALIX	<u>9MG</u>	<u>N203634</u>	<u>001</u>	Jan 14, 2013
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BUDESONIDE; FORMOTEROL FUMARATE DIHYDRATE

AEROSOL, METERED; INHALATION

SYMBICORT

	+	!	ASTRAZENECA	0.08MG/INH; 0.0045MG/INH	N021929	001	Jul 21, 2006
	+	!		0.16MG/INH; 0.0045MG/INH	N021929	002	Jul 21, 2006

BUMETANIDE

INJECTABLE; INJECTION

BUMETANIDE

<u>AP</u>		HOSPIRA	<u>0.25MG/ML</u>	<u>A074332</u>	<u>001</u>	Oct 31, 1994
<u>AP</u>	!	WEST-WARD PHARMS	<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>		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>		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>		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

PRESCRIPTION DRUG PRODUCT LIST

BUMETANIDE

TABLET; ORAL

BUMETANIDE

<u>AB</u>		<u>1MG</u>	<u>A202900 002</u>	Apr 30, 2018
<u>AB</u>		<u>2MG</u>	<u>A202900 003</u>	Apr 30, 2018
<u>BUMEX</u>				
<u>AB</u>	+	VALIDUS PHARMS	<u>0.5MG</u>	<u>N018225 002</u> Feb 28, 1983
<u>AB</u>	+		<u>1MG</u>	<u>N018225 001</u> Feb 28, 1983
<u>AB</u>	+		<u>2MG</u>	<u>N018225 003</u> Jun 14, 1985

BUPIVACAINE

INJECTABLE, LIPOSOMAL; INJECTION

EXPAREL

+	!	PACIRA PHARMS INC	133MG/10ML (13.3MG/ML)	N022496 001 Oct 28, 2011
+	!		266MG/20ML (13.3MG/ML)	N022496 002 Oct 28, 2011

BUPIVACAINE HYDROCHLORIDE

INJECTABLE; INJECTION

BUPIVACAINE HYDROCHLORIDE

<u>AP</u>		AUROBINDO PHARMA LTD	<u>0.25%</u>	<u>A207183 001</u> May 13, 2016
<u>AP</u>			<u>0.5%</u>	<u>A207183 002</u> May 13, 2016
<u>AP</u>		HOSPIRA	<u>0.25%</u>	<u>A070583 001</u> Feb 17, 1987
<u>AP</u>			<u>0.25%</u>	<u>A070586 001</u> Mar 03, 1987
<u>AP</u>			<u>0.25%</u>	<u>A070590 001</u> Feb 17, 1987
<u>AP</u>			<u>0.25%</u>	<u>N018053 002</u>
<u>AP</u>			<u>0.5%</u>	<u>A070584 001</u> Feb 17, 1986
<u>AP</u>			<u>0.5%</u>	<u>A070597 001</u> Mar 03, 1987
<u>AP</u>			<u>0.5%</u>	<u>A070609 001</u> Mar 03, 1987
<u>AP</u>			<u>0.5%</u>	<u>N018053 001</u>
<u>AP</u>			<u>0.75%</u>	<u>A070585 001</u> Mar 03, 1987
<u>AP</u>			<u>0.75%</u>	<u>N018053 003</u>
<u>BUPIVACAINE HYDROCHLORIDE PRESERVATIVE FREE</u>				
<u>AP</u>		AUROBINDO PHARMA LTD	<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>MARCAINE HYDROCHLORIDE</u>				
<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>		BAXTER HLTHCARE CORP	<u>0.75%</u>	<u>A207266 001</u> Jul 25, 2016
<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
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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
	!		<u>0.25%; 0.005MG/ML</u>	<u>A071165 001</u> Jun 16, 1988
			<u>0.25%; 0.005MG/ML</u>	<u>A071167 001</u> Jun 16, 1988

PRESCRIPTION DRUG PRODUCT LIST

BUPIVACAINE HYDROCHLORIDE; EPINEPHRINE BITARTRATE
 INJECTABLE; INJECTION

MARCAINE HYDROCHLORIDE W/ EPINEPHRINE

AP	+ !	HOSPIRA	<u>0.25%;0.0091MG/ML</u>	<u>N016964 004</u>
AP	+ !		<u>0.5%;0.0091MG/ML</u>	<u>N016964 008</u>

MARCAINE HYDROCHLORIDE W/ EPINEPHRINE PRESERVATIVE FREE

AP	+ !	HOSPIRA	<u>0.25%;0.0091MG/ML</u>	<u>N016964 013</u>
AP	+ !		<u>0.5%;0.0091MG/ML</u>	<u>N016964 007</u>
AP	+ !		<u>0.75%;0.0091MG/ML</u>	<u>N016964 010</u>

SENSORCAINE

AP		FRESENIUS KABI USA	<u>0.25%;0.0091MG/ML</u>	<u>A070966 001</u>	Oct 13, 1987
AP			<u>0.25%;0.0091MG/ML</u>	<u>A070967 001</u>	Oct 13, 1987
AP			<u>0.5%;0.0091MG/ML</u>	<u>A070968 001</u>	Oct 13, 1987
AP			<u>0.5%;0.0091MG/ML</u>	<u>N018304 004</u>	Sep 02, 1983
AP			<u>0.75%;0.0091MG/ML</u>	<u>N018304 005</u>	Sep 02, 1983
		BUPIVACAINE HYDROCHLORIDE AND EPINEPHRINE			
		! SEPTODONT	0.5%;0.0091MG/ML	A077250 001	Sep 27, 2006

BUPRENORPHINE

FILM, EXTENDED RELEASE; TRANSDERMAL

BUPRENORPHINE

AB		WATSON LABS TEVA	<u>5MCG/HR</u>	<u>A204937 001</u>	Nov 20, 2018
AB			<u>10MCG/HR</u>	<u>A204937 002</u>	Nov 20, 2018
AB			<u>15MCG/HR</u>	<u>A204937 003</u>	Nov 20, 2018
AB			<u>20MCG/HR</u>	<u>A204937 004</u>	Nov 20, 2018

BUTRANS

AB	+	PURDUE PHARMA LP	<u>5MCG/HR</u>	<u>N021306 001</u>	Jun 30, 2010
AB	+		<u>10MCG/HR</u>	<u>N021306 002</u>	Jun 30, 2010
AB	+		<u>15MCG/HR</u>	<u>N021306 004</u>	Jul 25, 2013
AB	+ !		<u>20MCG/HR</u>	<u>N021306 003</u>	Jun 30, 2010
	+		7.5MCG/HR	N021306 005	Jun 30, 2014

SOLUTION, EXTENDED RELEASE; SUBCUTANEOUS

SUBLOCADE

	+	INDIVIOR INC	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

IMPLANT; IMPLANTATION

PROBUPHINE

	+ !	TITAN PHARMS	EQ 80MG BASE/IMPLANT	N204442 001	May 26, 2016
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INJECTABLE; INJECTION

BUPRENEX

AP	+ !	INDIVIOR INC	<u>EQ 0.3MG BASE/ML</u>	<u>N018401 001</u>
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BUPRENORPHINE HYDROCHLORIDE

AP		AM REGENT	<u>EQ 0.3MG BASE/ML</u>	<u>A078331 001</u>	Mar 27, 2007
AP		HOSPIRA	<u>EQ 0.3MG BASE/ML</u>	<u>A074137 001</u>	Jun 03, 1996
AP		PAR STERILE PRODUCTS	<u>EQ 0.3MG BASE/ML</u>	<u>A206586 001</u>	Jul 28, 2015
AP		WEST-WARD PHARMS	<u>EQ 0.3MG BASE/ML</u>	<u>A076931 001</u>	Mar 02, 2005

TABLET; SUBLINGUAL

BUPRENORPHINE HYDROCHLORIDE

AB		ACTAVIS ELIZABETH	<u>EQ 2MG BASE</u>	<u>A090819 001</u>	Feb 19, 2015
AB			<u>EQ 8MG BASE</u>	<u>A090819 002</u>	Feb 19, 2015
AB		CASI PHARMS INC	<u>EQ 2MG BASE</u>	<u>A090279 001</u>	Jun 10, 2015
AB			<u>EQ 8MG BASE</u>	<u>A090279 002</u>	Jun 10, 2015
AB		ETHYPHARM	<u>EQ 2MG BASE</u>	<u>A090622 001</u>	Sep 24, 2010
AB			<u>EQ 8MG BASE</u>	<u>A090622 002</u>	Sep 24, 2010
AB		HIKMA	<u>EQ 2MG BASE</u>	<u>A078633 001</u>	Oct 08, 2009
AB	!		<u>EQ 8MG BASE</u>	<u>A078633 002</u>	Oct 08, 2009
AB		RHODES PHARMS	<u>EQ 2MG BASE</u>	<u>A207276 001</u>	Mar 27, 2017
AB			<u>EQ 8MG BASE</u>	<u>A207276 002</u>	Mar 27, 2017
AB		SUN PHARM	<u>EQ 2MG BASE</u>	<u>A201760 001</u>	Jan 29, 2016
AB			<u>EQ 8MG BASE</u>	<u>A201760 002</u>	Jan 29, 2016

PRESCRIPTION DRUG PRODUCT LIST

3-66 (of 453)

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;BUCCAL, SUBLINGUAL

BUPRENORPHINE HYDROCHLORIDE AND NALOXONE HYDROCHLORIDE

AB	ALVOGEN PINE BROOK	<u>EQ 2MG BASE;EQ 0.5MG BASE</u>	A205954	001	Jan 24, 2019
AB		<u>EQ 4MG BASE;EQ 1MG BASE</u>	A205954	002	Jan 24, 2019
AB		<u>EQ 8MG BASE;EQ 2MG BASE</u>	A205954	003	Jan 24, 2019
AB		<u>EQ 12MG BASE;EQ 3MG BASE</u>	A205954	004	Jan 24, 2019
AB	DR REDDYS LABS SA	<u>EQ 2MG BASE;EQ 0.5MG BASE</u>	A205299	001	Jun 14, 2018
AB		<u>EQ 4MG BASE;EQ 1MG BASE</u>	A205806	001	Jun 14, 2018
AB		<u>EQ 8MG BASE;EQ 2MG BASE</u>	A205299	002	Jun 14, 2018
AB		<u>EQ 12MG BASE;EQ 3MG BASE</u>	A205806	002	Jun 14, 2018
AB	MYLAN TECHNOLOGIES	<u>EQ 8MG BASE;EQ 2MG BASE</u>	A207607	001	Jun 14, 2018
AB		<u>EQ 12MG BASE;EQ 3MG BASE</u>	A207607	002	Jun 14, 2018

SUBOXONE

AB	+	INDIVIOR INC	<u>EQ 2MG BASE;EQ 0.5MG BASE</u>	N022410	001	Aug 30, 2010
AB	+		<u>EQ 4MG BASE;EQ 1MG BASE</u>	N022410	003	Aug 10, 2012
AB	+		<u>EQ 8MG BASE;EQ 2MG BASE</u>	N022410	002	Aug 30, 2010
AB	+	!	<u>EQ 12MG BASE;EQ 3MG BASE</u>	N022410	004	Aug 10, 2012

TABLET;SUBLINGUAL

BUPRENORPHINE HYDROCHLORIDE AND NALOXONE HYDROCHLORIDE

AB	ACTAVIS ELIZABETH	<u>EQ 2MG BASE;EQ 0.5MG BASE</u>	A091422	001	Feb 22, 2013
AB	!	<u>EQ 8MG BASE;EQ 2MG BASE</u>	A091422	002	Feb 22, 2013
AB	AMNEAL PHARMS	<u>EQ 2MG BASE;EQ 0.5MG BASE</u>	A203136	001	Feb 22, 2013
AB		<u>EQ 8MG BASE;EQ 2MG BASE</u>	A203136	002	Feb 22, 2013
AB	ETHYPHARM USA CORP	<u>EQ 2MG BASE;EQ 0.5MG BASE</u>	A204431	001	Oct 16, 2015
AB		<u>EQ 8MG BASE;EQ 2MG BASE</u>	A204431	002	Oct 16, 2015
AB	HIKMA	<u>EQ 2MG BASE;EQ 0.5MG BASE</u>	A203326	001	Jun 27, 2014
AB		<u>EQ 8MG BASE;EQ 2MG BASE</u>	A203326	002	Jun 27, 2014
AB	LANNETT CO INC	<u>EQ 2MG BASE;EQ 0.5MG BASE</u>	A205022	001	Sep 19, 2016
AB		<u>EQ 8MG BASE;EQ 2MG BASE</u>	A205022	002	Sep 19, 2016
AB	SPECGX LLC	<u>EQ 2MG BASE;EQ 0.5MG BASE</u>	A207000	001	Dec 13, 2017
AB		<u>EQ 8MG BASE;EQ 2MG BASE</u>	A207000	002	Dec 13, 2017
AB	SUN PHARM	<u>EQ 2MG BASE;EQ 0.5MG BASE</u>	A201633	001	Aug 05, 2016
AB		<u>EQ 8MG BASE;EQ 2MG BASE</u>	A201633	002	Aug 05, 2016

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

+	VALEANT PHARMS NORTH	174MG	N022108	001	Apr 23, 2008
+		348MG	N022108	002	Apr 23, 2008
+	!	522MG	N022108	003	Apr 23, 2008

BUPROPION HYDROCHLORIDE

TABLET;ORAL

BUPROPION HYDROCHLORIDE

AB	ALEMBIC PHARMS LTD	75MG	A203013	001	Jun 08, 2018
AB		100MG	A203013	002	Jun 08, 2018
AB	APNAR PHARMA LP	75MG	A075584	001	Feb 07, 2000
AB		100MG	A075584	002	Feb 07, 2000
AB	APOTEX INC	75MG	A076143	001	Jan 17, 2006
AB	!	100MG	A076143	002	Jan 17, 2006
AB	CADILA PHARMS LTD	75MG	A208606	001	Jan 16, 2020
AB		100MG	A208606	002	Jan 16, 2020
AB	HERITAGE PHARMA	75MG	A206975	001	Aug 19, 2016
AB		100MG	A206975	002	Aug 19, 2016
AB	INVAGEN PHARMS	75MG	A207389	001	Sep 18, 2017
AB		100MG	A207389	002	Sep 18, 2017
AB	MYLAN	75MG	A075491	001	Apr 17, 2000
AB		100MG	A075491	002	Apr 17, 2000

PRESCRIPTION DRUG PRODUCT LIST

BUPROPION HYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL

BUPROPION HYDROCHLORIDE

<u>AB1</u>	ACTAVIS LABS FL INC	<u>100MG</u>	<u>A079095 001</u>	Mar 24, 2009
<u>AB1</u>		<u>150MG</u>	<u>A079095 002</u>	Mar 24, 2009
<u>AB1</u>		<u>200MG</u>	<u>A079095 003</u>	Mar 24, 2009
<u>AB1</u>	ANCHEN PHARMS	<u>100MG</u>	<u>A091459 001</u>	Jun 09, 2011
<u>AB1</u>		<u>150MG</u>	<u>A091459 002</u>	Jun 09, 2011
<u>AB1</u>		<u>200MG</u>	<u>A091459 003</u>	Jun 09, 2011
<u>AB1</u>	IMPAX LABS	<u>100MG</u>	<u>A075913 001</u>	Jan 28, 2004
<u>AB1</u>		<u>150MG</u>	<u>A075913 002</u>	Mar 22, 2004
<u>AB1</u>		<u>200MG</u>	<u>A076711 001</u>	Dec 03, 2004
<u>AB1</u>	INVAGEN PHARMS	<u>100MG</u>	<u>A206674 001</u>	Feb 09, 2016
<u>AB1</u>		<u>150MG</u>	<u>A206674 002</u>	Feb 09, 2016
<u>AB1</u>		<u>200MG</u>	<u>A206674 003</u>	Feb 09, 2016
<u>AB1</u>	PRINSTON INC	<u>100MG</u>	<u>A202304 001</u>	May 26, 2015
<u>AB1</u>		<u>150MG</u>	<u>A202304 002</u>	May 26, 2015
<u>AB1</u>		<u>200MG</u>	<u>A202304 003</u>	May 26, 2015
<u>AB1</u>	SANDOZ	<u>100MG</u>	<u>A075932 001</u>	Nov 25, 2003
<u>AB1</u>		<u>150MG</u>	<u>A075932 002</u>	Mar 22, 2004
<u>AB1</u>		<u>200MG</u>	<u>A075932 003</u>	Jun 22, 2005
<u>AB1</u>	SCIEGEN PHARMS INC	<u>100MG</u>	<u>A205794 001</u>	Mar 01, 2016
<u>AB1</u>		<u>150MG</u>	<u>A205794 002</u>	Mar 01, 2016
<u>AB1</u>		<u>200MG</u>	<u>A205794 003</u>	Mar 01, 2016
<u>AB1</u>	SUN PHARM	<u>100MG</u>	<u>A078866 001</u>	Apr 06, 2010
<u>AB1</u>		<u>150MG</u>	<u>A078866 002</u>	Apr 06, 2010
<u>AB1</u>		<u>200MG</u>	<u>A078866 003</u>	Apr 06, 2010
<u>AB1</u>	TORRENT	<u>100MG</u>	<u>A203969 001</u>	Oct 31, 2014
<u>AB1</u>		<u>150MG</u>	<u>A203969 002</u>	Oct 31, 2014
<u>AB1</u>		<u>200MG</u>	<u>A203969 003</u>	Oct 31, 2014
<u>AB1</u>	WATSON LABS INC	<u>100MG</u>	<u>A077455 001</u>	Jul 19, 2010
<u>AB1</u>		<u>150MG</u>	<u>A077455 002</u>	Mar 12, 2008
<u>AB1</u>		<u>200MG</u>	<u>A077455 003</u>	Jul 19, 2010

WELLBUTRIN SR

<u>AB1</u>	+	GLAXOSMITHKLINE	<u>100MG</u>	<u>N020358 002</u>	Oct 04, 1996
<u>AB1</u>	+		<u>150MG</u>	<u>N020358 003</u>	Oct 04, 1996
<u>AB1</u>	+	!	<u>200MG</u>	<u>N020358 004</u>	Jun 14, 2002

BUPROPION HYDROCHLORIDE

<u>AB2</u>	!	ACTAVIS LABS FL INC	<u>150MG</u>	<u>A079094 001</u>	Mar 24, 2009
<u>AB2</u>		ANCHEN PHARMS	<u>150MG</u>	<u>A091520 001</u>	Jun 09, 2011
<u>AB2</u>		IMPAX LABS	<u>150MG</u>	<u>A075914 001</u>	May 27, 2004
<u>AB2</u>		SANDOZ INC	<u>150MG</u>	<u>A077475 001</u>	Mar 12, 2008
<u>AB2</u>		SCIEGEN PHARMS INC	<u>150MG</u>	<u>A206122 001</u>	Aug 17, 2016
<u>AB3</u>		ACCORD HLTHCARE	<u>150MG</u>	<u>A210497 001</u>	Oct 31, 2018
<u>AB3</u>			<u>300MG</u>	<u>A210497 002</u>	Oct 31, 2018
<u>AB3</u>		ACTAVIS LABS FL INC	<u>150MG</u>	<u>A077715 001</u>	Nov 26, 2008
<u>AB3</u>		ANBISON LAB	<u>150MG</u>	<u>A207224 001</u>	Jun 30, 2017
<u>AB3</u>			<u>300MG</u>	<u>A207224 002</u>	Jun 30, 2017
<u>AB3</u>		ANCHEN PHARMS	<u>150MG</u>	<u>A077284 001</u>	Dec 14, 2006
<u>AB3</u>			<u>300MG</u>	<u>A077284 002</u>	Dec 14, 2006
<u>AB3</u>		GRAVITI PHARMS	<u>150MG</u>	<u>A211020 001</u>	Jan 28, 2019
<u>AB3</u>			<u>300MG</u>	<u>A211020 002</u>	Jan 28, 2019
<u>AB3</u>		IMPAX LABS	<u>150MG</u>	<u>A077415 001</u>	Nov 26, 2008
<u>AB3</u>		INVAGEN PHARMS	<u>150MG</u>	<u>A206556 001</u>	Aug 26, 2016
<u>AB3</u>			<u>300MG</u>	<u>A206556 002</u>	Aug 26, 2016
<u>AB3</u>		LUPIN LTD	<u>150MG</u>	<u>A090693 001</u>	Apr 06, 2017
<u>AB3</u>			<u>300MG</u>	<u>A090693 002</u>	Apr 06, 2017
<u>AB3</u>		SCIEGEN PHARMS INC	<u>150MG</u>	<u>A207479 001</u>	Apr 12, 2017
<u>AB3</u>			<u>300MG</u>	<u>A207479 002</u>	Apr 12, 2017
<u>AB3</u>		SINOTHERAPEUTICS INC	<u>150MG</u>	<u>A208652 001</u>	Aug 21, 2017
<u>AB3</u>			<u>300MG</u>	<u>A208652 002</u>	Aug 21, 2017
<u>AB3</u>		TWI PHARMS	<u>150MG</u>	<u>A210081 001</u>	Nov 03, 2017
<u>AB3</u>			<u>300MG</u>	<u>A210081 002</u>	Nov 03, 2017
<u>AB3</u>		WATSON LABS INC	<u>150MG</u>	<u>A077285 001</u>	Nov 26, 2008
<u>AB3</u>			<u>300MG</u>	<u>A077285 002</u>	Aug 15, 2008
<u>AB3</u>		WOCKHARDT LTD	<u>150MG</u>	<u>A202189 001</u>	Nov 21, 2012
<u>AB3</u>		YICHANG HUMANWELL	<u>150MG</u>	<u>A210015 001</u>	Jun 14, 2018
<u>AB3</u>			<u>300MG</u>	<u>A210015 002</u>	Jun 14, 2018
<u>AB3</u>		ZHEJIANG JUTAI PHARM	<u>300MG</u>	<u>A211200 001</u>	Sep 05, 2019
<u>AB3</u>		ZYDUS PHARMS	<u>150MG</u>	<u>A201567 002</u>	Jul 23, 2018
<u>AB3</u>			<u>300MG</u>	<u>A201567 001</u>	Jan 17, 2014

PRESCRIPTION DRUG PRODUCT LIST

BUPROPION HYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL

WELLBUTRIN XL

AB3	+	VALEANT INTL	150MG	N021515	001	Aug 28, 2003
AB3	+	!	300MG	N021515	002	Aug 28, 2003
		FORFIVO XL				
	+	ALVOGEN	450MG	N022497	001	Nov 10, 2011

BUPROPION HYDROCHLORIDE; NALTREXONE HYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL

CONTRAVE

+	NALPROPION	90MG;8MG	N200063	001	Sep 10, 2014
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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		AMNEAL PHARMS CO	5MG	A208829	001	May 24, 2017
AB			7.5MG	A208829	002	May 24, 2017
AB			10MG	A208829	003	May 24, 2017
AB			15MG	A208829	004	May 24, 2017
AB			30MG	A208829	005	May 24, 2017
AB		AUROBINDO PHARMA LTD	5MG	A078246	001	Feb 27, 2009
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	A075467	002	Mar 28, 2001
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	30MG	A078302	001	Dec 17, 2007
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
AB	!		15MG	A075022	003	Feb 28, 2002
AB			30MG	A075022	004	Mar 25, 2004
AB		UNICHEM LABS LTD	5MG	A210907	001	Nov 14, 2019
AB			10MG	A210907	002	Nov 14, 2019
AB			15MG	A210907	003	Nov 14, 2019
AB			30MG	A210907	004	Nov 14, 2019
AB		YILING PHARM LTD	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		ZYDUS PHARMS	5MG	A078888	001	Feb 07, 2014
AB			10MG	A078888	002	Feb 07, 2014

PRESCRIPTION DRUG PRODUCT LIST

BUSPIRONE HYDROCHLORIDE

TABLET; ORAL

BUSPIRONE HYDROCHLORIDE

AB		15MG	A078888 003	Feb 07, 2014
AB		30MG	A078888 004	Feb 07, 2014

BUSULFAN

INJECTABLE; INJECTION

BUSULFAN

AP	ACCORD HLTHCARE INC	6MG/ML	A210148 001	Feb 22, 2019
AP	ACTAVIS LLC	6MG/ML	A205139 001	Dec 08, 2017
AP	AMNEAL	6MG/ML	A209580 001	Dec 18, 2017
AP	APOTEX	6MG/ML	A210448 001	May 07, 2019
AP	ATHENEX INC	6MG/ML	A205106 001	Sep 21, 2018
AP	HOSPIRA INC	6MG/ML	A205672 001	Jul 31, 2018
AP	LUITPOLD	6MG/ML	A202259 001	Dec 22, 2015
AP	NEXUS PHARMS	6MG/ML	A207794 001	Jan 14, 2019
AP	PHARMASCIENCE INC	6MG/ML	A207050 001	Mar 24, 2017
AP	SHILPA MEDICARE LTD	6MG/ML	A210931 001	Apr 18, 2019

BUSULFEX

AP	+!	OTSUKA PHARM	6MG/ML	N020954 001	Feb 04, 1999
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MYLERAN

AP	MYLAN INSTITUTIONAL	6MG/ML	A208536 001	Nov 20, 2017
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TABLET; ORAL

MYLERAN

+!	ASPEN GLOBAL	2MG	N009386	001
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BUTABARBITAL SODIUM

TABLET; ORAL

BUTISOL SODIUM

+!	MYLAN SPECIALITY LP	30MG	N000793	004
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BUTENAFINE HYDROCHLORIDE

CREAM; TOPICAL

MENTAX

+!	MYLAN	1%	N020524	001	Oct 18, 1996
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BUTOCONAZOLE NITRATE

CREAM; VAGINAL

GYNAZOLE-1

!	PERRIGO ISRAEL	2%	A200923	001	May 18, 2012
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BUTORPHANOL TARTRATE

INJECTABLE; INJECTION

BUTORPHANOL TARTRATE

AP	HIKMA FARMACEUTICA	1MG/ML	A078400 001	May 01, 2009
AP		2MG/ML	A078400 002	May 01, 2009
AP	WEST-WARD PHARMS	2MG/ML	A075046 001	Aug 12, 1998

INT

BUTORPHANOL TARTRATE PRESERVATIVE FREE

AP	!	HOSPIRA	1MG/ML	A074626 001	Jan 23, 1997
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AP	!		2MG/ML	A074626 002	Jan 23, 1997
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AP		WEST-WARD PHARMS	1MG/ML	A075045 001	Aug 12, 1998
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INT

AP			2MG/ML	A075045 002	Aug 12, 1998
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SPRAY, METERED; NASAL

BUTORPHANOL TARTRATE

AB	APOTEX INC	1MG/SPRAY	A075499 001	Dec 04, 2002
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AB	HIKMA	1MG/SPRAY	A075824 001	Mar 12, 2002
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AB	!	MYLAN	1MG/SPRAY	A075759 001	Aug 08, 2001
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CABAZITAXEL

SOLUTION; INTRAVENOUS

JEVTANA KIT

+!	SANOFI AVENTIS US	60MG/1.5ML (40MG/ML)	N201023	001	Jun 17, 2010
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CABERGOLINE

TABLET; ORAL

CABERGOLINE

AB	INGENUS PHARMS LLC	0.5MG	A204735 001	Aug 01, 2018
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AB	IVAX SUB TEVA	0.5MG	A077750 001	Mar 07, 2007
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PHARMS

AB	!	PAR PHARM	0.5MG	A076310 001	Dec 29, 2005
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PRESCRIPTION DRUG PRODUCT LIST

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

CAFICITAP +! HIKMAEQ 30MG BASE/3ML (EQ 10MG BASE/ML)N020793 001 Sep 21, 1999CAFFEINE CITRATEAP AM REGENTEQ 30MG BASE/3ML (EQ 10MG BASE/ML)A077906 001 May 15, 2007AP AUROBINDO PHARMA
LTDEQ 30MG BASE/3ML (EQ 10MG BASE/ML)A205013 001 Sep 22, 2015AP EXELA PHARMA
SCIENCEEQ 30MG BASE/3ML (EQ 10MG BASE/ML)A077233 001 Sep 21, 2006AP FRESENIUS KABI USAEQ 30MG BASE/3ML (EQ 10MG BASE/ML)A077997 001 Jul 20, 2007AP MICRO LABSEQ 30MG BASE/3ML (EQ 10MG BASE/ML)A207400 001 Dec 14, 2017AP SAGENT PHARMSEQ 30MG BASE/3ML (EQ 10MG BASE/ML)A090827 001 Aug 29, 2012AP SUN PHARMEQ 30MG BASE/3ML (EQ 10MG BASE/ML)A090077 001 Sep 30, 2009

SOLUTION; ORAL

CAFICITAA +! HIKMAEQ 30MG BASE/3ML (EQ 10MG BASE/ML)N020793 002 Apr 12, 2000CAFFEINE CITRATEAA EXELA PHARMA SCS
LLCEQ 30MG BASE/3ML (EQ 10MG BASE/ML)A077304 001 Sep 21, 2006AA FRESENIUS KABI USAEQ 30MG BASE/3ML (EQ 10MG BASE/ML)A078002 001 Jan 31, 2008AA MICRO LABSEQ 30MG BASE/3ML (EQ 10MG BASE/ML)A213202 001 Dec 16, 2019AA SAGENT PHARMSEQ 30MG BASE/3ML (EQ 10MG BASE/ML)A091102 001 Aug 29, 2012AA SUN PHARMEQ 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

CAFERGOTAA ! SANDOZ100MG; 1MGA084294 001ERGOTAMINE TARTRATE AND CAFFEINEAA HIKMA INTL PHARMS100MG; 1MGA040510 001 Sep 17, 2004AA MIKART100MG; 1MGA040590 001 Sep 16, 2005CALCIFEDIOL

CAPSULE, EXTENDED RELEASE; ORAL

RAYALDEE

+! OPKO IRELAND GLOBAL 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 PHARMS0.005%A205772 001 Jun 09, 2015AB TOLMAR0.005%A200935 001 May 30, 2012DOVONEXAB +! LEO PHARMA AS0.005%N020554 001 Jul 22, 1996

OINTMENT; TOPICAL

CALCIPOTRIENE

! GLENMARK PHARMS INC 0.005%

A090633 001 Mar 24, 2010

SOLUTION; TOPICAL

CALCIPOTRIENEAT ACP NIMBLE0.005%A078468 001 Mar 24, 2011AT FOUGERA PHARMS0.005%A078305 001 May 06, 2008AT HI TECH PHARMA0.005%A077579 001 Nov 19, 2009AT NOVEL LABS INC0.005%A207163 001 Dec 26, 2017AT ! TOLMAR0.005%A077029 001 Nov 20, 2009

PRESCRIPTION DRUG PRODUCT LIST

CALCITONIN SALMON

INJECTABLE; INJECTION

MIACALCIN

+	!	MYLAN IRELAND LTD	200 IU/ML	N017808	002	Mar 29, 1991
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SPRAY, METERED; NASAL

CALCITONIN-SALMON

<u>AB</u>	!	APOTEX INC	<u>200 IU/SPRAY</u>	<u>A076396</u>	<u>001</u>	Nov 17, 2008
<u>AB</u>		PAR PHARM	<u>200 IU/SPRAY</u>	<u>A076979</u>	<u>001</u>	Jun 08, 2009

CALCITRIOL

CAPSULE; ORAL

CALCITRIOL

<u>AB</u>		AMNEAL PHARMS	<u>0.25MCG</u>	<u>A203289</u>	<u>002</u>	Jun 14, 2017
<u>AB</u>			<u>0.5MCG</u>	<u>A203289</u>	<u>001</u>	Jun 14, 2017
<u>AB</u>		BIONPHARMA INC	<u>0.25MCG</u>	<u>A091174</u>	<u>001</u>	May 24, 2013
<u>AB</u>			<u>0.5MCG</u>	<u>A091174</u>	<u>002</u>	May 24, 2013
<u>AB</u>		HIKMA	<u>0.25MCG</u>	<u>A076917</u>	<u>001</u>	Mar 27, 2006
<u>AB</u>		STRIDES PHARMA	<u>0.25MCG</u>	<u>A091356</u>	<u>001</u>	Dec 12, 2014
<u>AB</u>			<u>0.5MCG</u>	<u>A091356</u>	<u>002</u>	Dec 12, 2014
<u>AB</u>		TEVA	<u>0.25MCG</u>	<u>A075765</u>	<u>001</u>	Oct 12, 2001
<u>AB</u>			<u>0.5MCG</u>	<u>A075765</u>	<u>002</u>	Oct 12, 2001

ROCALTROL

<u>AB</u>	+	VALIDUS PHARMS	<u>0.25MCG</u>	<u>N018044</u>	<u>001</u>	
<u>AB</u>	+	!	<u>0.5MCG</u>	<u>N018044</u>	<u>002</u>	

INJECTABLE; INJECTION

CALCITRIOL

<u>AP</u>	!	AKORN	<u>0.001MG/ML</u>	<u>A078066</u>	<u>001</u>	Jan 29, 2008
<u>AP</u>		GLAND PHARMA LTD	<u>0.001MG/ML</u>	<u>A211030</u>	<u>001</u>	Feb 03, 2020

OINTMENT; TOPICAL

VECTICAL

+	!	GALDERMA LABS LP	3MCG/GM	N022087	001	Jan 23, 2009
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SOLUTION; ORAL

CALCITRIOL

<u>AA</u>		ELYSIUM	<u>1MCG/ML</u>	<u>A209798</u>	<u>001</u>	Nov 21, 2018
<u>AA</u>		HIKMA	<u>1MCG/ML</u>	<u>A076242</u>	<u>001</u>	Jul 18, 2003

ROCALTROL

<u>AA</u>	+	!	VALIDUS PHARMS	<u>1MCG/ML</u>	<u>N021068</u>	<u>001</u>	Nov 20, 1998
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CALCIUM ACETATE

CAPSULE; ORAL

CALCIUM ACETATE

<u>AB</u>		AMNEAL PHARMS	<u>667MG</u>	<u>A201658</u>	<u>001</u>	Oct 06, 2014
<u>AB</u>		CHARTWELL RX	<u>667MG</u>	<u>A091312</u>	<u>001</u>	Jun 01, 2012
<u>AB</u>		HERITAGE PHARMS INC	<u>667MG</u>	<u>A202315</u>	<u>001</u>	Jun 29, 2015
<u>AB</u>		HIKMA	<u>667MG</u>	<u>A077728</u>	<u>001</u>	Feb 26, 2008
<u>AB</u>		INVAGEN PHARMS	<u>667MG</u>	<u>A203135</u>	<u>001</u>	Feb 07, 2013
<u>AB</u>		LUPIN LTD	<u>667MG</u>	<u>A202127</u>	<u>001</u>	Jul 09, 2015
<u>AB</u>		NOSTRUM LABS INC	<u>667MG</u>	<u>A203179</u>	<u>001</u>	Oct 26, 2015
<u>AB</u>		SUVEN LIFE	<u>667MG</u>	<u>A211038</u>	<u>001</u>	Feb 21, 2020

PHOSLO GELCAPS

<u>AB</u>	+	!	FRESENIUS MEDCL	<u>667MG</u>	<u>N021160</u>	<u>003</u>	Apr 02, 2001
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SOLUTION; ORAL

PHOSLYRA

+	!	FRESENIUS MEDCL	667MG/5ML	N022581	001	Apr 18, 2011
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TABLET; ORAL

CALCIUM ACETATE

<u>AB</u>		CHARTWELL MOLECULAR	<u>667MG</u>	<u>A202420</u>	<u>001</u>	Feb 05, 2013
<u>AB</u>		HERITAGE PHARMS INC	<u>667MG</u>	<u>A202885</u>	<u>001</u>	Jan 22, 2015
<u>AB</u>	!	PADDOCK LLC	<u>667MG</u>	<u>A091561</u>	<u>001</u>	Apr 13, 2011

CALCIUM CHLORIDE

INJECTABLE; INJECTION

CALCIUM CHLORIDE 10%

<u>AP</u>		AM REGENT	<u>100MG/ML</u>	<u>A209088</u>	<u>001</u>	Jul 27, 2017
<u>AP</u>		INTL MEDICATION SYS	<u>100MG/ML</u>	<u>A203477</u>	<u>001</u>	May 09, 2018
<u>AP</u>		MEDEFIL INC	<u>100MG/ML</u>	<u>A211553</u>	<u>001</u>	May 01, 2019

CALCIUM CHLORIDE 10% IN PLASTIC CONTAINER

<u>AP</u>	+	!	HOSPIRA	<u>100MG/ML</u>	<u>N021117</u>	<u>001</u>	Jan 28, 2000
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PRESCRIPTION DRUG PRODUCT LIST

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
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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
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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
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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
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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
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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
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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
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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
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CALCIUM CHLORIDE; DEXTROSE; MAGNESIUM CHLORIDE; SODIUM CHLORIDE; SODIUM LACTATE

SOLUTION; INTRAPERITONEAL

DELFLX W/ DEXTROSE 1.5% IN PLASTIC CONTAINER

AT	FRESENIUS MEDCL	25.7MG/100ML; 1.5GM/100ML; 15.2MG/100ML; 5 67MG/100ML; 392MG/100ML	N018883 001	Nov 30, 1984
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DELFLX W/ DEXTROSE 1.5% LOW MAGNESIUM IN PLASTIC CONTAINER

AT	FRESENIUS MEDCL	25.7MG/100ML; 1.5GM/100ML; 5.08MG/100ML; 5 38MG/100ML; 448MG/100ML	N018883 004	Nov 30, 1984
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DELFLX W/ DEXTROSE 1.5% LOW MAGNESIUM LOW CALCIUM IN PLASTIC CONTAINER

AT	FRESENIUS MEDCL	18.4MG/100ML; 1.5GM/100ML; 5.08MG/100ML; 5 38MG/100ML; 448MG/100ML	N020171 001	Aug 19, 1992
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DELFLX W/ DEXTROSE 2.5% IN PLASTIC CONTAINER

AT	FRESENIUS MEDCL	25.7MG/100ML; 2.5GM/100ML; 15.2MG/100ML; 5 67MG/100ML; 392MG/100ML	N018883 002	Nov 30, 1984
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DELFLX W/ DEXTROSE 2.5% LOW MAGNESIUM IN PLASTIC CONTAINER

AT	FRESENIUS MEDCL	25.7MG/100ML; 2.5GM/100ML; 5.08MG/100ML; 5 38MG/100ML; 448MG/100ML	N018883 005	Nov 30, 1984
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DELFLX W/ DEXTROSE 2.5% LOW MAGNESIUM LOW CALCIUM IN PLASTIC CONTAINER

AT	FRESENIUS MEDCL	18.4MG/100ML; 2.5GM/100ML; 5.08MG/100ML; 5 38MG/100ML; 448MG/100ML	N020171 002	Aug 19, 1992
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DELFLX W/ DEXTROSE 4.25% IN PLASTIC CONTAINER

AT	FRESENIUS MEDCL	25.7MG/100ML; 4.25GM/100ML; 15.2MG/100ML; 5 567MG/100ML; 392MG/100ML	N018883 003	Nov 30, 1984
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DELFLX W/ DEXTROSE 4.25% LOW MAGNESIUM IN PLASTIC CONTAINER

AT	FRESENIUS MEDCL	25.7MG/100ML; 4.25GM/100ML; 5.08MG/100ML; 5 538MG/100ML; 448MG/100ML	N018883 006	Nov 30, 1984
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DELFLX W/ DEXTROSE 4.25% LOW MAGNESIUM LOW CALCIUM IN PLASTIC CONTAINER

AT	FRESENIUS MEDCL	18.4MG/100ML; 4.25GM/100ML; 5.08MG/100ML; 5 538MG/100ML; 448MG/100ML	N020171 003	Aug 19, 1992
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DIANEAL LOW CALCIUM W/ DEXTROSE 1.5% IN PLASTIC CONTAINER

AT	+ BAXTER HLTHCARE	18.3MG/100ML; 1.5GM/100ML; 5.08MG/100ML; 5 38MG/100ML; 448MG/100ML	N020183 001	Dec 04, 1992
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DIANEAL LOW CALCIUM W/ DEXTROSE 2.5% IN PLASTIC CONTAINER

AT	+ BAXTER HLTHCARE	18.3MG/100ML; 2.5GM/100ML; 5.08MG/100ML; 5 38MG/100ML; 448MG/100ML	N020183 002	Dec 04, 1992
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DIANEAL LOW CALCIUM W/ DEXTROSE 4.25% IN PLASTIC CONTAINER

AT	+ BAXTER HLTHCARE	18.3MG/100ML; 4.25GM/100ML; 5.08MG/100ML; 5 538MG/100ML; 448MG/100ML	N020183 004	Dec 04, 1992
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PRESCRIPTION DRUG PRODUCT LIST

CALCIUM CHLORIDE; DEXTROSE; MAGNESIUM CHLORIDE; SODIUM CHLORIDE; SODIUM LACTATE
 SOLUTION; INTRAPERITONEAL

<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>N017512 012</u>	<u>Jan 10, 1989</u>
			<u>38MG/100ML;448MG/100ML</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>N017512 013</u>	<u>Jul 11, 1990</u>
			<u>38MG/100ML;448MG/100ML</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>N017512 014</u>	<u>Jul 11, 1990</u>
			<u>38MG/100ML;448MG/100ML</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>N017512 015</u>	<u>Jul 11, 1990</u>
			<u>538MG/100ML;448MG/100ML</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>N017512 004</u>	
			<u>38MG/100ML;448MG/100ML</u>		
<u>AT</u>	<u>+</u>		<u>25.7MG/100ML;1.5GM/100ML;5.08MG/100ML;5</u>	<u>N020163 001</u>	<u>Dec 04, 1992</u>
			<u>38MG/100ML;448MG/100ML</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>N017512 005</u>	
			<u>38MG/100ML;448MG/100ML</u>		
<u>AT</u>	<u>+</u>		<u>25.7MG/100ML;2.5GM/100ML;5.08MG/100ML;5</u>	<u>N020163 002</u>	<u>Dec 04, 1992</u>
			<u>38MG/100ML;448MG/100ML</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>N017512 006</u>	
			<u>538MG/100ML;448MG/100ML</u>		
<u>AT</u>	<u>+</u>		<u>25.7MG/100ML;4.25GM/100ML;5.08MG/100ML;</u>	<u>N020163 003</u>	<u>Dec 04, 1992</u>
			<u>538MG/100ML;448MG/100ML</u>		

CALCIUM CHLORIDE; DEXTROSE; MAGNESIUM SULFATE; POTASSIUM CHLORIDE; SODIUM BICARBONATE; SODIUM
CHLORIDE; SODIUM PHOSPHATE, DIBASIC, HEPTAHYDRATE

<u>INJECTABLE; INTRATHECAL</u>					
<u>ELLIOTTS B SOLUTION</u>					
	<u>+</u>	<u>LUKARE MEDICAL LLC</u>	<u>0.2MG/ML;0.8MG/ML;0.3MG/ML;0.3MG/ML;1.9</u>	<u>N020577 001</u>	<u>Sep 27, 1996</u>
			<u>MG/ML;7.3MG/ML;0.2MG/ML</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>N017608 001</u>	
			<u>00ML;310MG/100ML</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>N019634 003</u>	<u>Feb 24, 1988</u>
			<u>00ML;310MG/100ML</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>N016679 001</u>	
			<u>00ML;310MG/100ML</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>N019367 006</u>	<u>Apr 05, 1985</u>
			<u>100ML;310MG/100ML</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>N019367 004</u>	<u>Apr 05, 1985</u>
			<u>100ML;310MG/100ML</u>		
<u>AP</u>			<u>20MG/100ML;5GM/100ML;328MG/100ML;600MG/</u>	<u>N019367 005</u>	<u>Apr 05, 1985</u>
			<u>100ML;310MG/100ML</u>		
<u>AP</u>	<u>+</u>	<u>ICU MEDICAL INC</u>	<u>20MG/100ML;5GM/100ML;179MG/100ML;600MG/</u>	<u>N019685 002</u>	<u>Oct 17, 1988</u>
			<u>100ML;310MG/100ML</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>N019367 007</u>	<u>Apr 05, 1985</u>
			<u>100ML;310MG/100ML</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>N019367 008</u>	<u>Apr 05, 1985</u>
			<u>100ML;310MG/100ML</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;300MG</u>	<u>N019634 001</u>	<u>Feb 24, 1988</u>
			<u>/100ML;160MG/100ML</u>		
<u>POTASSIUM CHLORIDE 10MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER</u>					
		<u>BAXTER HLTHCARE</u>	<u>20MG/100ML;5GM/100ML;105MG/100ML;600MG/</u>	<u>N019367 002</u>	<u>Apr 05, 1985</u>
			<u>100ML;310MG/100ML</u>		
			<u>20MG/100ML;5GM/100ML;179MG/100ML;600MG/</u>	<u>N019367 003</u>	<u>Apr 05, 1985</u>
			<u>100ML;310MG/100ML</u>		
<u>POTASSIUM CHLORIDE 5MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER</u>					
		<u>BAXTER HLTHCARE</u>	<u>20MG/100ML;5GM/100ML;105MG/100ML;600MG/</u>	<u>N019367 001</u>	<u>Apr 05, 1985</u>
			<u>100ML;310MG/100ML</u>		

PRESCRIPTION DRUG PRODUCT LIST

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 AKORN 0.48MG/ML; 0.3MG/ML; 0.75MG/ML; 3.9MG/ML; 6.4MG/ML; 1.7MG/ML **A075503 001** Sep 27, 2006

AT B BRAUN 0.48MG/ML; 0.3MG/ML; 0.75MG/ML; 3.9MG/ML; 6.4MG/ML; 1.7MG/ML **A091387 001** Feb 03, 2010

BSS

AT +! ALCON 0.48MG/ML; 0.3MG/ML; 0.75MG/ML; 3.9MG/ML; 6.4MG/ML; 1.7MG/ML **N020742 001** Dec 10, 1997

CALCIUM CHLORIDE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE; SODIUM PHOSPHATE

INJECTABLE; INJECTION

PHOXILLUM B22K 4/0 IN PLASTIC CONTAINER

+! BAXTER HLTHCARE N/A/1000ML; 3.05GM/1000ML; 0.314GM/1000ML N207026 002 Jan 13, 2015
CORP ; 2.21GM/1000ML; 6.95GM/1000ML; 0.187GM/1000ML (5000ML)

PHOXILLUM BK 4/2.5 IN PLASTIC CONTAINER

+! BAXTER HLTHCARE 3.68GM/1000ML; 3.05GM/1000ML; 0.314GM/1000ML N207026 001 Jan 13, 2015
CORP OML ; 3.09GM/1000ML; 6.34GM/1000ML; 0.187GM/1000ML

CALCIUM CHLORIDE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM CHLORIDE

SOLUTION; PERFUSION, CARDIAC

CARDIOPLEGIC IN PLASTIC CONTAINER

AT BAXTER HLTHCARE 17.6MG/100ML; 325.3MG/100ML; 119.3MG/100ML; 643MG/100ML **A075323 001** Apr 21, 2000

PLEGISOL IN PLASTIC CONTAINER

AT +! HOSPIRA 17.6MG/100ML; 325.3MG/100ML; 119.3MG/100ML; 643MG/100ML **N018608 001** Feb 26, 1982

CALCIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM CHLORIDE

INJECTABLE; INJECTION

RINGER'S IN PLASTIC CONTAINER

AP B BRAUN 33MG/100ML; 30MG/100ML; 860MG/100ML **N020002 001** Apr 17, 1992

AP BAXTER HLTHCARE 33MG/100ML; 30MG/100ML; 860MG/100ML **N016693 001**

AP ICU MEDICAL INC 33MG/100ML; 30MG/100ML; 860MG/100ML **N018251 001**

SOLUTION; IRRIGATION

RINGER'S IN PLASTIC CONTAINER

AT B BRAUN 33MG/100ML; 30MG/100ML; 860MG/100ML **N018156 001**

AT BAXTER HLTHCARE 33MG/100ML; 30MG/100ML; 860MG/100ML **N018495 001** Feb 19, 1982

AT ICU MEDICAL INC 33MG/100ML; 30MG/100ML; 860MG/100ML **N017635 001**

CALCIUM 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/100ML **N019632 001** Feb 29, 1988

AP +! BAXTER HLTHCARE 20MG/100ML; 30MG/100ML; 600MG/100ML; 310MG/100ML **N016682 001**

AP FRESENIUS KABI USA 20MG/100ML; 30MG/100ML; 600MG/100ML; 310MG/100ML **A209338 001** Jan 28, 2019

AP ICU MEDICAL INC 20MG/100ML; 30MG/100ML; 600MG/100ML; 310MG/100ML **N017641 001**

SOLUTION; IRRIGATION

LACTATED RINGER'S IN PLASTIC CONTAINER

AT +! B BRAUN 20MG/100ML; 30MG/100ML; 600MG/100ML; 310MG/100ML **N018681 001** Dec 27, 1982

AT BAXTER HLTHCARE 20MG/100ML; 30MG/100ML; 600MG/100ML; 310MG/100ML **N018494 001** Feb 19, 1982

AT +! 20MG/100ML; 30MG/100ML; 600MG/100ML; 310MG/100ML **N018921 001** Apr 03, 1984

AT +! ICU MEDICAL INC 20MG/100ML; 30MG/100ML; 600MG/100ML; 310MG/100ML **N019416 001** Jan 17, 1986

PRESCRIPTION DRUG PRODUCT LISTCALCIUM GLUCONATE

SOLUTION; INTRAVENOUS

CALCIUM GLUCONATE

+	!	FRESENIUS KABI USA	1GM/10ML (100MG/ML)	N208418	001	Jun 15, 2017
+	!		5GM/50ML (100MG/ML)	N208418	002	Jun 15, 2017
+	!		10GM/100ML (100MG/ML)	N208418	003	Jun 15, 2017
CALCIUM GLUCONATE IN SODIUM CHLORIDE						
+	!	HQ SPCLT PHARMA	1GM/50ML (20MG/ML)	N210906	001	Oct 29, 2018
+	!		2GM/100ML (20MG/ML)	N210906	002	Oct 29, 2018

CALFACTANT

SUSPENSION; INTRATRACHEAL

INFASURF PRESERVATIVE FREE

+	!	ONY	35MG/ML	N020521	001	Jul 01, 1998
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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

AB	+	ANI PHARMS INC	4MG	N020838	001	Jun 04, 1998
AB	+		8MG	N020838	002	Jun 04, 1998
AB	+		16MG	N020838	003	Jun 04, 1998
AB	+	!	32MG	N020838	004	Jun 04, 1998

CANDESARTAN CILEXETIL

AB		ALEMBIC PHARMS LTD	4MG	A210302	001	Dec 04, 2018
AB			8MG	A210302	002	Dec 04, 2018
AB			16MG	A210302	003	Dec 04, 2018
AB			32MG	A209119	001	Jun 20, 2017
AB		MACLEODS PHARMS LTD	4MG	A203813	001	Dec 05, 2016
AB			8MG	A203813	002	Dec 05, 2016
AB			16MG	A203813	003	Dec 05, 2016
AB			32MG	A203813	004	Dec 05, 2016
AB		MYLAN	4MG	A078702	001	May 03, 2013
AB			8MG	A078702	002	May 03, 2013
AB			16MG	A078702	003	May 03, 2013
AB			32MG	A078702	004	May 03, 2013
AB		ZYDUS PHARMS	4MG	A091390	001	Aug 23, 2017
AB			8MG	A091390	002	Aug 23, 2017
AB			16MG	A091390	003	Aug 23, 2017
AB			32MG	A091390	004	Aug 23, 2017

CANDESARTAN CILEXETIL; HYDROCHLOROTHIAZIDE

TABLET; ORAL

ATACAND HCT

AB	+	ANI PHARMS INC	16MG; 12.5MG	N021093	001	Sep 05, 2000
AB	+		32MG; 12.5MG	N021093	002	Sep 05, 2000
AB	+	!	32MG; 25MG	N021093	003	May 16, 2008

CANDESARTAN CILEXETIL AND HYDROCHLOROTHIAZIDE

AB		DR REDDYS LABS LTD	16MG; 12.5MG	A202965	001	Jun 03, 2013
AB			32MG; 12.5MG	A202965	002	Jun 03, 2013
AB			32MG; 25MG	A202965	003	Jun 03, 2013
AB		MACLEODS PHARMS LTD	16MG; 12.5MG	A204100	001	Feb 27, 2015
AB			32MG; 12.5MG	A204100	002	Feb 27, 2015
AB			32MG; 25MG	A204100	003	Feb 27, 2015
AB		MYLAN	16MG; 12.5MG	A090704	001	Dec 04, 2012
AB			32MG; 12.5MG	A090704	002	Dec 04, 2012

PRESCRIPTION DRUG PRODUCT LIST

CANDESARTAN CILEXETIL; HYDROCHLOROTHIAZIDE

TABLET; ORAL

CANDESARTAN CILEXETIL AND HYDROCHLOROTHIAZIDE

AB		<u>32MG;25MG</u>	<u>A090704 003</u>	Dec 04, 2012
AB	PRINSTON INC	<u>16MG;12.5MG</u>	<u>A207455 001</u>	Apr 11, 2018
AB		<u>32MG;12.5MG</u>	<u>A207455 002</u>	Apr 11, 2018
AB		<u>32MG;25MG</u>	<u>A207455 003</u>	Apr 11, 2018
AB	ZYDUS PHARMS	<u>16MG;12.5MG</u>	<u>A203466 001</u>	Nov 27, 2017
AB		<u>32MG;12.5MG</u>	<u>A203466 002</u>	Nov 27, 2017
AB		<u>32MG;25MG</u>	<u>A203466 003</u>	Nov 27, 2017

CANGRELOR

POWDER; INTRAVENOUS

KENGREAL

+! CHIESI USA INC 50MG/VIAL N204958 001 Jun 22, 2015

CANNABIDIOL

SOLUTION; ORAL

EPIDIOLEX

+! GW RES LTD 100MG/ML N210365 001 Sep 28, 2018

CAPECITABINE

TABLET; ORAL

CAPECITABINE

AB	ACCORD HLTHCARE	<u>150MG</u>	<u>A202593 001</u>	Apr 23, 2015
AB		<u>500MG</u>	<u>A202593 002</u>	Apr 23, 2015
AB	ALKEM LABS LTD	<u>150MG</u>	<u>A207652 001</u>	Nov 24, 2017
AB		<u>500MG</u>	<u>A207652 002</u>	Nov 24, 2017
AB	AMNEAL PHARMS	<u>150MG</u>	<u>A204741 001</u>	Feb 28, 2017
AB		<u>500MG</u>	<u>A204741 002</u>	Feb 28, 2017
AB	EUGIA PHARMA	<u>150MG</u>	<u>A210604 001</u>	Apr 17, 2018
AB		<u>500MG</u>	<u>A210604 002</u>	Apr 17, 2018
AB	HIKMA	<u>150MG</u>	<u>A200483 001</u>	Jul 14, 2016
AB		<u>500MG</u>	<u>A200483 002</u>	Jul 14, 2016
AB	MSN	<u>150MG</u>	<u>A209365 001</u>	Jul 02, 2018
AB		<u>500MG</u>	<u>A209365 002</u>	Jul 02, 2018
AB	MYLAN	<u>150MG</u>	<u>A090943 001</u>	Aug 08, 2014
AB		<u>500MG</u>	<u>A090943 002</u>	Aug 08, 2014
AB	SHILPA MEDICARE LTD	<u>150MG</u>	<u>A207456 001</u>	Dec 12, 2016
AB		<u>500MG</u>	<u>A207456 002</u>	Dec 12, 2016
AB	SUN PHARM	<u>150MG</u>	<u>A204668 001</u>	Jun 21, 2019
AB		<u>500MG</u>	<u>A204668 002</u>	Jun 21, 2019
AB	TEVA PHARMS USA	<u>150MG</u>	<u>A091649 001</u>	Sep 16, 2013
AB		<u>500MG</u>	<u>A091649 002</u>	Sep 16, 2013

XELODA

AB	+ HOFFMANN LA ROCHE	<u>150MG</u>	<u>N020896 001</u>	Apr 30, 1998
AB	+!	<u>500MG</u>	<u>N020896 002</u>	Apr 30, 1998

CAPREOMYCIN SULFATE

INJECTABLE; INJECTION

CAPASTAT SULFATEAP +! AKORN EQ 1GM BASE/VIAL N050095 001CAPREOMYCIN SULFATE

AP	HISUN PHARM HANGZHOU	<u>EQ 1GM BASE/VIAL</u>	<u>A204796 001</u>	Oct 18, 2018
AP	MYLAN LABS LTD	<u>EQ 1GM BASE/VIAL</u>	<u>A202634 001</u>	Nov 27, 2017

CAPSAICIN

PATCH; TOPICAL

QUTENZA

+! AVERITAS 8% N022395 001 Nov 16, 2009

CAPTOPRIL

TABLET; ORAL

CAPTOPRIL

AB	AJANTA PHARMA LTD	<u>12.5MG</u>	<u>A212809 001</u>	Dec 13, 2019
AB		<u>25MG</u>	<u>A212809 002</u>	Dec 13, 2019
AB		<u>50MG</u>	<u>A212809 003</u>	Dec 13, 2019
AB		<u>100MG</u>	<u>A212809 004</u>	Dec 13, 2019
AB	BOSCOGEN	<u>12.5MG</u>	<u>A074677 004</u>	May 30, 1997
AB		<u>25MG</u>	<u>A074677 002</u>	May 30, 1997
AB		<u>50MG</u>	<u>A074677 001</u>	May 30, 1997
AB		<u>100MG</u>	<u>A074677 003</u>	May 30, 1997
AB	HIKMA INTL PHARMS	<u>12.5MG</u>	<u>A074505 001</u>	Feb 13, 1996
AB		<u>25MG</u>	<u>A074505 002</u>	Feb 13, 1996
AB		<u>50MG</u>	<u>A074505 003</u>	Feb 13, 1996

PRESCRIPTION DRUG PRODUCT LIST

CAPTOPRIL

TABLET; ORAL

CAPTOPRIL

AB		100MG	A074505 004	Feb 13, 1996
AB	MYLAN	12.5MG	A074434 001	Feb 13, 1996
AB		25MG	A074434 002	Feb 13, 1996
AB		50MG	A074434 003	Feb 13, 1996
AB	!	100MG	A074434 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	SETON PHARMS	12.5MG	A212223 001	Oct 30, 2019
AB		25MG	A212223 002	Oct 30, 2019
AB		50MG	A212223 003	Oct 30, 2019
AB		100MG	A212223 004	Oct 30, 2019
AB	TEVA	12.5MG	A074322 001	Feb 13, 1996
AB		25MG	A074322 002	Feb 13, 1996
AB		50MG	A074322 003	Feb 13, 1996
AB		100MG	A074322 004	Feb 13, 1996
AB	WATSON LABS	12.5MG	A074386 001	May 23, 1996
AB		25MG	A074386 002	May 23, 1996
AB		50MG	A074386 003	May 23, 1996
AB		100MG	A074386 004	May 23, 1996
AB	WOCKHARDT LTD	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

	MYLAN	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	
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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	TARO	100MG	A201106 001	Jun 21, 2013
AB		200MG	A201106 002	Jun 21, 2013
AB		300MG	A201106 003	Jun 21, 2013
AB	TEVA PHARMS	100MG	A078592 001	Sep 20, 2012
AB		200MG	A078592 002	Sep 20, 2012
AB		300MG	A078592 003	Sep 20, 2012

CARBATROL

AB	+	SHIRE DEV LLC	100MG	N020712 003	Sep 30, 1997
AB	+		200MG	N020712 001	Sep 30, 1997
AB	+	!	300MG	N020712 002	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

AB	WOCKHARDT BIO AG	100MG/5ML	A075714 001	Jun 05, 2002
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TEGRETOL

AB	+	NOVARTIS	100MG/5ML	N018927 001	Dec 18, 1987
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TERIL

AB	TARO	100MG/5ML	A076729 001	Sep 20, 2004
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TABLET; ORAL

CARBAMAZEPINE

AB	APOTEX INC	200MG	A075948 001	Feb 27, 2002
AB	TARO	200MG	A074649 001	Oct 03, 1996
AB	TORRENT PHARMS	200MG	A077272 002	Dec 07, 2005

PRESCRIPTION DRUG PRODUCT LIST

CARBAMAZEPINE

TABLET;ORAL

EPITOL

AB	TEVA	200MG	A070541 001	Sep 17, 1986
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TEGRETOL

AB	+!	NOVARTIS	200MG	N016608 001
		CARBAMAZEPINE		
		TORRENT PHARMS	100MG	A077272 001 Dec 07, 2005
			300MG	A077272 003 Dec 07, 2005
			400MG	A077272 004 Dec 07, 2005

TABLET, CHEWABLE;ORAL

CARBAMAZEPINE

AB	TARO PHARM INDS	100MG	A075687 001	Oct 24, 2000
AB	TORRENT PHARMS	100MG	A075712 001	Jul 05, 2001

EPITOL

AB	TEVA	100MG	A073524 001	Jul 29, 1992
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TEGRETOL

AB	+!	NOVARTIS	100MG	N018281 001
		CARBAMAZEPINE		
		! TARO PHARM INDS	200MG	A075687 002 Jul 29, 2002

TABLET, EXTENDED RELEASE;ORAL

CARBAMAZEPINE

AB	TARO	100MG	A078115 001	Mar 31, 2009
AB		200MG	A078115 002	Mar 31, 2009
AB		400MG	A078115 003	Mar 31, 2009
AB	ZYDUS PHARMS	100MG	A205571 001	Feb 07, 2019
AB		200MG	A205571 002	Feb 07, 2019
AB		400MG	A205571 003	Feb 07, 2019

TEGRETOL-XR

AB	+	NOVARTIS	100MG	N020234 001	Mar 25, 1996
AB	+		200MG	N020234 002	Mar 25, 1996
AB	+!		400MG	N020234 003	Mar 25, 1996

CARBIDOPA

TABLET;ORAL

CARBIDOPA

AB	ALVOGEN	25MG	A204291 001	Jan 08, 2016
AB	ANI PHARMS INC	25MG	A203261 001	Mar 10, 2014
AB	AUROBINDO PHARMA LTD	25MG	A211055 001	Oct 21, 2019
AB	EDENBRIDGE PHARMS	25MG	A205304 001	Feb 17, 2016
AB	NOVEL LABS INC	25MG	A204763 001	Oct 20, 2017
AB	ZYDUS PHARMS	25MG	A209910 001	May 07, 2018

LODOSYN

AB	+!	ATON	25MG	N017830 001
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CARBIDOPA; ENTACAPONE; LEVODOPA

TABLET;ORAL

CARBIDOPA, LEVODOPA AND ENTACAPONE

AB	SUN PHARM	25MG;200MG;100MG	A079085 001	May 10, 2012	
AB		37.5MG;200MG;150MG	A079085 002	May 10, 2012	
	STALEVO 100				
AB	+	ORION PHARMA	25MG;200MG;100MG	N021485 002	Jun 11, 2003
	STALEVO 150				
AB	+	ORION PHARMA	37.5MG;200MG;150MG	N021485 003	Jun 11, 2003
	STALEVO 125				
	+	ORION PHARMA	31.25MG;200MG;125MG	N021485 006 Aug 29, 2008	
	STALEVO 200				
	+	ORION PHARMA	50MG;200MG;200MG	N021485 004 Aug 02, 2007	
	STALEVO 50				
	+	ORION PHARMA	12.5MG;200MG;50MG	N021485 001 Jun 11, 2003	
	STALEVO 75				
	+	ORION PHARMA	18.75MG;200MG;75MG	N021485 005 Aug 29, 2008	

CARBIDOPA; LEVODOPA

CAPSULE, EXTENDED RELEASE;ORAL

RYTARY

	+	IMPAX LABS INC	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 INC	4.63MG/ML;20MG/ML	N203952 001	Jan 09, 2015
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PRESCRIPTION DRUG PRODUCT LIST

CARBIDOPA; LEVODOPA

TABLET; ORAL

CARBIDOPA AND LEVODOPA

<u>AB</u>	ACTAVIS ELIZABETH	<u>10MG;100MG</u>	<u>A074260 001</u>	Sep 03, 1993
<u>AB</u>		<u>25MG;100MG</u>	<u>A074260 002</u>	Sep 03, 1993
<u>AB</u>		<u>25MG;250MG</u>	<u>A074260 003</u>	Sep 03, 1993
<u>AB</u>	APOTEX INC	<u>10MG;100MG</u>	<u>A077120 001</u>	Jun 02, 2008
<u>AB</u>		<u>25MG;100MG</u>	<u>A077120 002</u>	Jun 02, 2008
<u>AB</u>		<u>25MG;250MG</u>	<u>A077120 003</u>	Jun 02, 2008
<u>AB</u>	MAYNE PHARMA	<u>10MG;100MG</u>	<u>A073618 001</u>	Aug 28, 1992
<u>AB</u>		<u>25MG;100MG</u>	<u>A073589 001</u>	Aug 28, 1992
<u>AB</u>		<u>25MG;250MG</u>	<u>A073607 001</u>	Aug 28, 1992
<u>AB</u>	MYLAN	<u>10MG;100MG</u>	<u>A090324 001</u>	Sep 28, 2009
<u>AB</u>		<u>25MG;100MG</u>	<u>A090324 002</u>	Sep 28, 2009
<u>AB</u>		<u>25MG;250MG</u>	<u>A090324 003</u>	Sep 28, 2009
<u>AB</u>	SUN PHARM INDS	<u>10MG;100MG</u>	<u>A078536 001</u>	Oct 28, 2008
<u>AB</u>		<u>25MG;100MG</u>	<u>A078536 002</u>	Oct 28, 2008
<u>AB</u>		<u>25MG;250MG</u>	<u>A078536 003</u>	Oct 28, 2008

SINEMET

<u>AB</u>	+ MERCK SHARP DOHME	<u>10MG;100MG</u>	<u>N017555 001</u>	
<u>AB</u>	+	<u>25MG;100MG</u>	<u>N017555 003</u>	
<u>AB</u>	+!	<u>25MG;250MG</u>	<u>N017555 002</u>	

TABLET, EXTENDED RELEASE; ORAL

CARBIDOPA AND LEVODOPA

<u>AB</u>	ACCORD HLTHCARE	<u>25MG;100MG</u>	<u>A202323 001</u>	Feb 08, 2013
<u>AB</u>		<u>50MG;200MG</u>	<u>A202323 002</u>	Feb 08, 2013
<u>AB</u>	ALEMBIC PHARMS LTD	<u>25MG;100MG</u>	<u>A210341 001</u>	Jun 05, 2019
<u>AB</u>		<u>50MG;200MG</u>	<u>A210341 002</u>	Jun 05, 2019
<u>AB</u>	APOTEX	<u>25MG;100MG</u>	<u>A076212 001</u>	Jun 16, 2004
<u>AB</u>		<u>50MG;200MG</u>	<u>A076212 002</u>	Jun 16, 2004
<u>AB</u>	IMPAX LABS	<u>25MG;100MG</u>	<u>A076521 001</u>	May 14, 2004
<u>AB</u>		<u>50MG;200MG</u>	<u>A076521 002</u>	May 14, 2004
<u>AB</u>	MYLAN	<u>25MG;100MG</u>	<u>A075091 002</u>	Apr 21, 2000
<u>AB</u>		<u>50MG;200MG</u>	<u>A075091 001</u>	Sep 30, 1999
<u>AB</u>	SUN PHARM INDS	<u>25MG;100MG</u>	<u>A077828 001</u>	Aug 23, 2007
<u>AB</u>	!	<u>50MG;200MG</u>	<u>A077828 002</u>	Aug 23, 2007

TABLET, ORALLY DISINTEGRATING; ORAL

CARBIDOPA AND LEVODOPA

<u>AB</u>	MYLAN	<u>10MG;100MG</u>	<u>A078893 001</u>	Sep 18, 2008
<u>AB</u>		<u>25MG;100MG</u>	<u>A078893 002</u>	Sep 18, 2008
<u>AB</u>	!	<u>25MG;250MG</u>	<u>A078893 003</u>	Sep 18, 2008
<u>AB</u>	SUN PHARM	<u>10MG;100MG</u>	<u>A078690 001</u>	Jul 31, 2009
<u>AB</u>		<u>25MG;100MG</u>	<u>A078690 002</u>	Jul 31, 2009
<u>AB</u>		<u>25MG;250MG</u>	<u>A078690 003</u>	Jul 31, 2009

CARBINOXAMINE MALEATE

SOLUTION; ORAL

CARBINOXAMINE MALEATE

! MIKART 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

<u>AA</u>	INVAGEN PHARMS	<u>4MG</u>	<u>A090435 001</u>	Apr 15, 2010
<u>AA</u>	!	<u>4MG</u>	<u>A040442 001</u>	Mar 19, 2003
<u>AA</u>	MISSION PHARMACAL CO	<u>4MG</u>	<u>A090756 001</u>	May 27, 2011
	!	MIKART 6MG	A207484 001	May 31, 2016

CARBOPLATIN

INJECTABLE; INTRAVENOUS

CARBOPLATIN

<u>AP</u>	ACCORD HLTHCARE	<u>50MG/5ML (10MG/ML)</u>	<u>A206775 001</u>	Feb 09, 2017
<u>AP</u>		<u>150MG/15ML (10MG/ML)</u>	<u>A206775 002</u>	Feb 09, 2017
<u>AP</u>		<u>450MG/45ML (10MG/ML)</u>	<u>A206775 003</u>	Feb 09, 2017
<u>AP</u>		<u>600MG/60ML (10MG/ML)</u>	<u>A206775 004</u>	Feb 09, 2017
<u>AP</u>	AKORN	<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>	CIPLA LTD	<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

PRESCRIPTION DRUG PRODUCT LIST

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CARBOPLATIN

INJECTABLE; INTRAVENOUS

CARBOPLATIN

<u>AP</u>		<u>600MG/60ML (10MG/ML)</u>	<u>A077861 004</u>	Jan 18, 2007
<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>	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>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>	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>	INGENUS PHARMS LLC	<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>	MEITHEAL	<u>50MG/5ML (10MG/ML)</u>	<u>A077096 001</u>	Jun 14, 2005
<u>AP</u>		<u>150MG/15ML (10MG/ML)</u>	<u>A077096 002</u>	Jun 14, 2005
<u>AP</u>		<u>450MG/45ML (10MG/ML)</u>	<u>A077096 003</u>	Jun 14, 2005
<u>AP</u>		<u>600MG/60ML (10MG/ML)</u>	<u>A077096 004</u>	Jun 03, 2013
<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>	PLIVA LACHEMA	<u>50MG/5ML (10MG/ML)</u>	<u>A078631 001</u>	Dec 02, 2008
<u>AP</u>		<u>150MG/15ML (10MG/ML)</u>	<u>A078631 002</u>	Dec 02, 2008
<u>AP</u>		<u>450MG/45ML (10MG/ML)</u>	<u>A078631 003</u>	Dec 02, 2008
<u>AP</u>		<u>600MG/60ML (10MG/ML)</u>	<u>A078631 004</u>	Dec 02, 2008
<u>AP</u>	SANDOZ INC	<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>	! TEVA PHARMS USA	<u>50MG/5ML (10MG/ML)</u>	<u>A077139 001</u>	Sep 21, 2005
<u>AP</u>	!	<u>150MG/15ML (10MG/ML)</u>	<u>A077139 002</u>	Sep 21, 2005
<u>AP</u>	!	<u>450MG/45ML (10MG/ML)</u>	<u>A077139 003</u>	Sep 21, 2005
<u>AP</u>	!	<u>600MG/60ML (10MG/ML)</u>	<u>A077139 004</u>	Sep 21, 2005
<u>AP</u>	WEST-WARD PHARMS INT	<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

INJECTABLE; IV (INFUSION)

CARBOPLATIN

<u>AP</u>	GLAND PHARMA LTD	<u>50MG/5ML (10MG/ML)</u>	<u>A207324 001</u>	Feb 15, 2017
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CARBOPROST TROMETHAMINE

INJECTABLE; INJECTION

CARBOPROST TROMETHAMINE

<u>AP</u>	DR REDDYS LABS LTD	<u>EQ 0.25MG BASE/ML</u>	<u>A211941 001</u>	Jul 02, 2019
<u>AP</u>	<u>HEMABATE</u>			
<u>AP</u>	+! PHARMACIA AND UPJOHN	<u>EQ 0.25MG BASE/ML</u>	<u>N017989 001</u>	

CARFILZOMIB

POWDER; INTRAVENOUS

CARFILZOMIB

<u>AP</u>	APOTEX	<u>60MG/VIAL</u>	<u>A209425 001</u>	Mar 16, 2020
<u>AP</u>	DR REDDYS	<u>60MG/VIAL</u>	<u>A209422 001</u>	Sep 09, 2019
<u>AP</u>	<u>KYPROLIS</u>			
<u>AP</u>	+! ONYX THERAP	<u>60MG/VIAL</u>	<u>N202714 001</u>	Jul 20, 2012
	+	10MG/VIAL	N202714 003	Jun 07, 2018
	+	30MG/VIAL	N202714 002	Jun 03, 2016

PRESCRIPTION DRUG PRODUCT LIST

CARGLUMIC ACID

TABLET; ORAL

CARBAGLU

+! RECORDATI RARE 200MG N022562 001 Mar 18, 2010

CARIPRAZINE HYDROCHLORIDE

CAPSULE; ORAL

VRAYLAR

+ ALLERGAN 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

CARISOPRODOLAA ACCELRX LABS 350MG A040576 001 Jun 07, 2005AA ALLIED 350MG A040245 001 Sep 08, 1997AA AUROBINDO PHARMA 350MG A040792 001 Aug 06, 2009AA NATCO PHARMA LTD 350MG A090988 001 Oct 28, 2014AA NOVAST LABS 350MG A040823 001 Oct 22, 2008AA ORIENT PHARMA CO 350MG A205085 001 Oct 28, 2014AA LTDAA SCIEGEN PHARMS INC 350MG A203374 001 Jan 27, 2014AA STRIDES PHARMA 350MG A205513 002 Nov 12, 2015AA WATSON LABS 350MG A087499 001 Apr 20, 1982AA WILSHIRE PHARMS INC 350MG A205126 002 Jul 08, 2015SOMAAA + MYLAN SPECIALITY LP 350MG N011792 001CARISOPRODOLAB AUROBINDO PHARMA 250MG A040792 002 Nov 08, 2016AB NOSTRUM LABS INC 250MG A207237 001 May 11, 2017AB STRIDES PHARMA 250MG A205513 001 Nov 12, 2015AB WILSHIRE PHARMS INC 250MG A205126 001 Jul 08, 2015SOMAAB +! MYLAN SPECIALITY LP 250MG N011792 004 Sep 13, 2007CARMUSTINE

IMPLANT; INTRACRANIAL

GLIADEL

+! ARBOR PHARMS LLC 7.7MG N020637 001 Sep 23, 1996

INJECTABLE; INJECTION

BICNUAP +! EMCURE PHARMS LTD 100MG/VIAL N017422 001CARMUSTINEAP AMNEAL 100MG/VIAL A211229 001 Oct 16, 2018AP NAVINTA LLC 100MG/VIAL A210179 001 Sep 11, 2018AP STI PHARMA LLC 100MG/VIAL A209278 001 Apr 02, 2019CARTEOLOL HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

CARTEOLOL HYDROCHLORIDE

! SANDOZ INC 1% A075476 001 Jan 03, 2000

CARVEDILOL

TABLET; ORAL

CARVEDILOLAB AUROBINDO PHARMA 3.125MG A078332 001 Sep 05, 2007AB 6.25MG A078332 002 Sep 05, 2007AB 12.5MG A078332 003 Sep 05, 2007AB 25MG A078332 004 Sep 05, 2007AB BEXIMCO USA 3.125MG A078384 001 Sep 05, 2007AB 6.25MG A078384 002 Sep 05, 2007AB 12.5MG A078384 003 Sep 05, 2007AB 25MG A078384 004 Sep 05, 2007AB CHARTWELL MOLECULAR 3.125MG A077474 001 Sep 05, 2007AB 6.25MG A077474 002 Sep 05, 2007AB 12.5MG A077474 003 Sep 05, 2007AB 25MG A077474 004 Sep 05, 2007AB DR REDDYS LABS LTD 3.125MG A076649 001 Sep 05, 2007AB 6.25MG A076649 002 Sep 05, 2007AB 12.5MG A076649 003 Sep 05, 2007AB 25MG A076649 004 Sep 05, 2007AB GLENMARK GENERICS 3.125MG A078251 001 Sep 05, 2007AB 6.25MG A078251 002 Sep 05, 2007

PRESCRIPTION DRUG PRODUCT LIST

3-82 (of 453)

CARVEDILOL

TABLET; ORAL

CARVEDILOL

<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>	MYLAN	<u>3.125MG</u>	<u>A077316 001</u>	Sep 05, 2007
<u>AB</u>		<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>	SUN PHARM INDS LTD	<u>3.125MG</u>	<u>A076989 001</u>	Sep 05, 2007
<u>AB</u>		<u>6.25MG</u>	<u>A076989 002</u>	Sep 05, 2007
<u>AB</u>		<u>12.5MG</u>	<u>A076989 003</u>	Sep 05, 2007
<u>AB</u>		<u>25MG</u>	<u>A076989 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>	+	SMITHKLINE BEECHAM	<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

COREG CR

<u>AB</u>	+	SMITHKLINE BEECHAM	<u>10MG</u>	<u>N022012 001</u>	Oct 20, 2006
<u>AB</u>	+		<u>20MG</u>	<u>N022012 002</u>	Oct 20, 2006
<u>AB</u>	+	!	<u>40MG</u>	<u>N022012 003</u>	Oct 20, 2006
<u>AB</u>	+		<u>80MG</u>	<u>N022012 004</u>	Oct 20, 2006

CASPOFUNGIN ACETATE

POWDER; INTRAVENOUS

CANCIDAS

<u>AP</u>	+	!	MERCK	<u>50MG/VIAL</u>	<u>N021227 001</u>	Jan 26, 2001
<u>AP</u>	+	!		<u>70MG/VIAL</u>	<u>N021227 002</u>	Jan 26, 2001

CASPOFUNGIN ACETATE

<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 PHARMA LTD	<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>		JIANGSU HENGRUI MED	<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

PRESCRIPTION DRUG PRODUCT LIST

CASPOFUNGIN ACETATE

POWDER; INTRAVENOUS

CASPOFUNGIN ACETATE

<u>AP</u>	MYLAN LABS LTD	<u>50MG/VIAL</u>	<u>A207650 001</u>	Sep 29, 2017
<u>AP</u>		<u>70MG/VIAL</u>	<u>A207650 002</u>	Sep 29, 2017
<u>AP</u>	XELLIA PHARMS APS	<u>50MG/VIAL</u>	<u>A205923 001</u>	Jul 02, 2018
<u>AP</u>		<u>70MG/VIAL</u>	<u>A205923 002</u>	Jul 02, 2018

CEFACTOR

CAPSULE; ORAL

CEFACTOR

<u>AB</u>	HIKMA	<u>EQ 250MG BASE</u>	<u>A065350 001</u>	Apr 03, 2007
<u>AB</u>	!	<u>EQ 500MG BASE</u>	<u>A065350 002</u>	Apr 03, 2007
<u>AB</u>	YUNG SHIN PHARM	<u>EQ 250MG BASE</u>	<u>A065146 001</u>	Jan 22, 2004
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>A065146 002</u>	Jan 22, 2004

FOR SUSPENSION; ORAL

CEFACTOR

	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

CEFACTOR

	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>	HIKMA	<u>EQ 500MG BASE</u>	<u>A065311 001</u>	Feb 07, 2006
<u>AB</u>	LUPIN	<u>EQ 500MG BASE</u>	<u>A065392 001</u>	May 29, 2007
<u>AB</u>	ORCHID HLTHCARE	<u>EQ 500MG BASE</u>	<u>A065309 001</u>	Sep 18, 2006
<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>	HIKMA PHARMS	<u>EQ 250MG BASE/5ML</u>	<u>A091036 001</u>	Nov 28, 2012
<u>AB</u>		<u>EQ 500MG BASE/5ML</u>	<u>A091036 002</u>	Nov 28, 2012
<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
<u>AB</u>	ORCHID HLTHCARE	<u>EQ 250MG BASE/5ML</u>	<u>A065307 002</u>	Oct 16, 2006
<u>AB</u>		<u>EQ 500MG BASE/5ML</u>	<u>A065307 003</u>	Oct 16, 2006

TABLET; ORAL

CEFADROXIL

<u>AB</u>	HIKMA	<u>EQ 1GM BASE</u>	<u>A065260 001</u>	Mar 30, 2006
<u>AB</u>	ORCHID HLTHCARE	<u>EQ 1GM BASE</u>	<u>A065301 001</u>	Sep 18, 2006
	! 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

ANCEF IN PLASTIC CONTAINER

	! BAXTER HLTHCARE	EQ 20MG BASE/ML	A063002 002	Mar 28, 1991
	CEFAZOLIN AND DEXTROSE			
	+! B BRAUN	EQ 1GM BASE/VIAL	N050779 002	Jul 27, 2000
	+	EQ 2GM BASE/VIAL	N050779 003	Jan 13, 2012
	CEFAZOLIN SODIUM			
	! ACS DOBFAR	EQ 20GM BASE/VIAL	A065306 002	Aug 18, 2014
	! 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

SOLUTION; INTRAVENOUS

CEFAZOLIN IN PLASTIC CONTAINER

BAXTER HLTHCARE	EQ 2GM BASE/100ML (EQ 20MG BASE/ML)	N207131 001	Aug 07, 2015
CORP			

CEFDINIR

CAPSULE; ORAL

CEFDINIR

<u>AB</u>	AUROBINDO PHARMA	<u>300MG</u>	<u>A065434 001</u>	Jan 07, 2008
<u>AB</u>	LUPIN	<u>300MG</u>	<u>A065264 001</u>	May 19, 2006
<u>AB</u>	ORCHID HLTHCARE	<u>300MG</u>	<u>A065418 001</u>	Jul 18, 2007
<u>AB</u>	! SANDOZ	<u>300MG</u>	<u>A065330 001</u>	Apr 06, 2007
<u>AB</u>	TEVA PHARMS	<u>300MG</u>	<u>A065368 001</u>	May 09, 2007

FOR SUSPENSION; ORAL

CEFDINIR

<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>	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>	ORCHID HLTHCARE	<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>	SANDOZ	<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>	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>	QILU	<u>EQ 500MG BASE/VIAL</u>	<u>A203704 001</u>	Feb 01, 2016
<u>AP</u>		<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

MAXIPIME

<u>AP</u>	+! HOSPIRA INC	<u>EQ 500MG BASE/VIAL</u>	<u>N050679 001</u>	Jan 18, 1996
<u>AP</u>	+!	<u>EQ 1GM BASE/VIAL</u>	<u>N050679 002</u>	Jan 18, 1996
<u>AP</u>	+!	<u>EQ 2GM BASE/VIAL</u>	<u>N050679 003</u>	Jan 18, 1996

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 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
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CEFIDEROCOL SULFATE TOSYLATE

POWDER; INTRAVENOUS

FETROJA

+!	SHIONOGI INC	EQ 1GM BASE/VIAL	N209445 001	Nov 14, 2019
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CEFIXIME

CAPSULE; ORAL

CEFIXIME

<u>AB</u>	ALKEM LABS LTD	<u>400MG</u>	<u>A210574 001</u>	Oct 09, 2018
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SUPRAX

<u>AB</u>	+! LUPIN LTD	<u>400MG</u>	<u>N203195 001</u>	Jun 01, 2012
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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

PRESCRIPTION DRUG PRODUCT LISTCEFIXIMETABLET, CHEWABLE; ORAL
SUPRAX

LUPIN LTD	100MG	A065380 001	Oct 25, 2010
	150MG	A065380 002	Oct 25, 2010
!	200MG	A065380 003	Oct 25, 2010

CEFOTAXIME SODIUM

INJECTABLE; INJECTION

CEFOTAXIME

! 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

CEFOTETAN DISODIUM

INJECTABLE; INJECTION

CEFOTAN

AP + TELIGENT	EQ 1GM BASE/VIAL	N050588 001	Dec 27, 1985
AP +	EQ 2GM BASE/VIAL	N050588 002	Dec 27, 1985

CEFOTETAN

AP ! FRESENIUS KABI USA	EQ 1GM BASE/VIAL	A065374 001	Aug 09, 2007
AP !	EQ 2GM BASE/VIAL	A065374 002	Aug 09, 2007
AP HIKMA	EQ 1GM BASE/VIAL	A091031 001	Oct 26, 2011
AP	EQ 2GM BASE/VIAL	A091031 002	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

CEFOXITIN SODIUM

INJECTABLE; INJECTION

CEFOXITIN

AP ! ACS DOBFAR	EQ 1GM BASE/VIAL	A065414 001	Jun 12, 2009
AP !	EQ 2GM BASE/VIAL	A065414 002	Jun 12, 2009
AP !	EQ 10GM BASE/VIAL	A065415 001	May 19, 2010
AP HIKMA FARMACEUTICA	EQ 1GM BASE/VIAL	A065238 001	Mar 12, 2010
AP	EQ 2GM BASE/VIAL	A065238 002	Mar 12, 2010
AP	EQ 10GM BASE/VIAL	A065239 001	Mar 02, 2010
AP WEST-WARD PHARMS	EQ 1GM BASE/VIAL	A065051 001	Sep 11, 2000
INT			
AP	EQ 2GM BASE/VIAL	A065051 002	Sep 11, 2000
AP	EQ 10GM BASE/VIAL	A065050 001	Sep 11, 2000
<u>CEFOXITIN AND DEXTROSE IN DUPLEX CONTAINER</u>			
AP +! B BRAUN	EQ 1GM BASE/VIAL	N065214 001	Mar 10, 2006
AP +!	EQ 2GM BASE/VIAL	N065214 002	Mar 10, 2006

MEFOXIN IN PLASTIC CONTAINER

! MYLAN INSTITUTIONAL	EQ 20MG BASE/ML	A063182 001	Jan 25, 1993
!	EQ 40MG BASE/ML	A063182 002	Jan 25, 1993

POWDER; INTRAVENOUS

CEFOXITIN IN PLASTIC CONTAINER

SAMSON MEDCL	EQ 100GM BASE	A200938 001	Nov 16, 2015
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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

AB AUROBINDO PHARMA	EQ 100MG BASE	A065370 001	Jun 11, 2007
AB	EQ 200MG BASE	A065370 002	Jun 11, 2007
AB ORCHID HLTHCARE	EQ 100MG BASE	A065388 001	Nov 14, 2007
AB	EQ 200MG BASE	A065388 002	Nov 14, 2007
AB SANDOZ	EQ 100MG BASE	A065462 001	May 28, 2008
AB !	EQ 200MG BASE	A065462 002	May 28, 2008

CEFPROZIL

FOR SUSPENSION; ORAL

CEFPROZIL

AB APOTEX INC	125MG/5ML	A065351 001	Feb 29, 2012
AB	250MG/5ML	A065351 002	Feb 29, 2012
AB AUROBINDO PHARMA	125MG/5ML	A065381 001	Jan 30, 2007
AB	250MG/5ML	A065381 002	Jan 30, 2007
AB LUPIN	125MG/5ML	A065261 001	Dec 19, 2005

PRESCRIPTION DRUG PRODUCT LIST

CEFPROZIL

FOR SUSPENSION; ORAL

CEFPROZIL

<u>AB</u>	!	<u>250MG/5ML</u>	<u>A065261</u>	<u>002</u>	Dec 19, 2005
<u>AB</u>	ORCHID HLTHCARE	<u>125MG/5ML</u>	<u>A065284</u>	<u>002</u>	Dec 30, 2005
<u>AB</u>		<u>250MG/5ML</u>	<u>A065284</u>	<u>001</u>	Dec 30, 2005
<u>AB</u>	SANDOZ	<u>125MG/5ML</u>	<u>A065257</u>	<u>001</u>	Dec 08, 2005
<u>AB</u>		<u>250MG/5ML</u>	<u>A065257</u>	<u>002</u>	Dec 08, 2005
<u>AB</u>	TEVA PHARMS	<u>125MG/5ML</u>	<u>A065236</u>	<u>001</u>	Dec 08, 2005
<u>AB</u>		<u>250MG/5ML</u>	<u>A065236</u>	<u>002</u>	Dec 08, 2005

TABLET; ORAL

CEFPROZIL

<u>AB</u>	APOTEX INC	<u>250MG</u>	<u>A065327</u>	<u>001</u>	Mar 26, 2008
<u>AB</u>		<u>500MG</u>	<u>A065327</u>	<u>002</u>	Mar 26, 2008
<u>AB</u>	AUROBINDO PHARMA LTD	<u>250MG</u>	<u>A065340</u>	<u>001</u>	May 24, 2007
<u>AB</u>		<u>500MG</u>	<u>A065340</u>	<u>002</u>	May 24, 2007
<u>AB</u>	CASI PHARMS INC	<u>250MG</u>	<u>A065235</u>	<u>001</u>	Nov 14, 2005
<u>AB</u>		<u>500MG</u>	<u>A065235</u>	<u>002</u>	Nov 14, 2005
<u>AB</u>	LUPIN	<u>250MG</u>	<u>A065276</u>	<u>001</u>	Dec 08, 2005
<u>AB</u>	!	<u>500MG</u>	<u>A065276</u>	<u>002</u>	Dec 08, 2005
<u>AB</u>	ORCHID HLTHCARE	<u>250MG</u>	<u>A065267</u>	<u>001</u>	Dec 19, 2005
<u>AB</u>		<u>500MG</u>	<u>A065267</u>	<u>002</u>	Dec 19, 2005
<u>AB</u>	TEVA	<u>250MG</u>	<u>A065208</u>	<u>001</u>	Dec 06, 2005
<u>AB</u>		<u>500MG</u>	<u>A065208</u>	<u>002</u>	Dec 06, 2005
<u>AB</u>	WOCKHARDT	<u>250MG</u>	<u>A065428</u>	<u>001</u>	Jun 14, 2007
<u>AB</u>		<u>500MG</u>	<u>A065428</u>	<u>002</u>	Jun 14, 2007

CEFTAROLINE FOSAMIL

POWDER; INTRAVENOUS

TEFLARO

+	ALLERGAN	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</u>	<u>002</u>	Nov 20, 1985
<u>AP</u>		<u>2GM/VIAL</u>	<u>A062640</u>	<u>003</u>	Nov 20, 1985
<u>AP</u>		<u>6GM/VIAL</u>	<u>A062640</u>	<u>004</u>	Feb 03, 1992
<u>AP</u>	WOCKHARDT	<u>1GM/VIAL</u>	<u>A065196</u>	<u>001</u>	Oct 15, 2008

FORTAZ

<u>AP</u>	+!	TELGENT	<u>500MG/VIAL</u>	<u>N050578</u>	<u>001</u>	Jul 19, 1985
<u>AP</u>	+	!	<u>1GM/VIAL</u>	<u>N050578</u>	<u>002</u>	Jul 19, 1985
<u>AP</u>	+	!	<u>2GM/VIAL</u>	<u>N050578</u>	<u>003</u>	Jul 19, 1985
<u>AP</u>	+	!	<u>6GM/VIAL</u>	<u>N050578</u>	<u>004</u>	Jul 19, 1985

TAZICEF

<u>AP</u>	HOSPIRA	<u>500MG/VIAL</u>	<u>A062662</u>	<u>001</u>	Mar 06, 1986
<u>AP</u>		<u>1GM/VIAL</u>	<u>A062662</u>	<u>002</u>	Mar 06, 1986
<u>AP</u>		<u>1GM/VIAL</u>	<u>A064032</u>	<u>001</u>	Oct 31, 1993
<u>AP</u>		<u>2GM/VIAL</u>	<u>A062662</u>	<u>003</u>	Mar 06, 1986
<u>AP</u>		<u>2GM/VIAL</u>	<u>A064032</u>	<u>002</u>	Oct 31, 1993
<u>AP</u>		<u>6GM/VIAL</u>	<u>A062662</u>	<u>004</u>	Mar 06, 1986

CEFTAZIDIME IN DEXTROSE CONTAINER

+	B BRAUN	EQ 1GM BASE	N050823	001	Jun 13, 2011
+	!	EQ 2GM BASE	N050823	002	Jun 13, 2011

CEFTOLOZANE SULFATE; TAZOBACTAM SODIUM

POWDER; INTRAVENOUS

ZERBAXA

+	CUBIST PHARMS LLC	EQ 1GM BASE/VIAL;EQ 0.5GM BASE/VIAL	N206829	001	Dec 19, 2014
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CEFTRIAZONE SODIUM

INJECTABLE; INJECTION

CEFTRIAZONE

<u>AP</u>	ACS DOBFAR	<u>EQ 500MG BASE/VIAL</u>	<u>A065329</u>	<u>001</u>	Jul 24, 2008	
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>A065329</u>	<u>002</u>	Jul 24, 2008	
<u>AP</u>		<u>EQ 2GM BASE/VIAL</u>	<u>A065329</u>	<u>003</u>	Jul 24, 2008	
<u>AP</u>		<u>EQ 10GM BASE/VIAL</u>	<u>A065328</u>	<u>001</u>	Jul 24, 2008	
<u>AP</u>	QILU	<u>EQ 10GM BASE/VIAL</u>	<u>A209218</u>	<u>001</u>	Oct 17, 2018	
<u>AP</u>	!	SANDOZ	<u>EQ 10GM BASE/VIAL</u>	<u>A065168</u>	<u>001</u>	May 17, 2005
<u>AP</u>	!	SANDOZ INC	<u>EQ 1GM BASE/VIAL</u>	<u>A065204</u>	<u>001</u>	May 03, 2005
<u>AP</u>	!		<u>EQ 2GM BASE/VIAL</u>	<u>A065204</u>	<u>002</u>	May 03, 2005
<u>AP</u>	WOCKHARDT	<u>EQ 1GM BASE/VIAL</u>	<u>A065180</u>	<u>001</u>	May 12, 2006	

PRESCRIPTION DRUG PRODUCT LIST

CEFTRIAXONE SODIUM

INJECTABLE; INJECTION

CEFTRIAXONE AND DEXTROSE IN DUPLEX CONTAINER

<u>AP</u>	<u>+</u> !	<u>B BRAUN</u>	<u>EQ 1GM BASE/VIAL</u>	<u>N050796 001</u>	Apr 20, 2005
<u>AP</u>	<u>+</u> !		<u>EQ 2GM BASE/VIAL</u>	<u>N050796 002</u>	Apr 20, 2005

CEFTRIAXONE SODIUM

<u>AP</u>		<u>ASTRAL</u>	<u>EQ 10GM BASE/VIAL</u>	<u>A091117 001</u>	Jan 20, 2017
<u>AP</u>		<u>HIKMA</u>	<u>EQ 10GM BASE/VIAL</u>	<u>A090701 001</u>	Oct 04, 2017

CEFTRIAXONE

		<u>SAMSON MEDCL</u>	<u>EQ 100GM BASE/VIAL</u>	<u>A090057 001</u>	Apr 25, 2014
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CEFTRIAXONE IN PLASTIC CONTAINER

<u>!</u>		<u>BAXTER HLTHCARE</u>	<u>EQ 20MG BASE/ML</u>	<u>A065224 001</u>	Aug 23, 2005
<u>!</u>			<u>EQ 40MG BASE/ML</u>	<u>A065224 002</u>	Aug 23, 2005

INJECTABLE; INTRAMUSCULAR, INTRAVENOUS

CEFTRIAXONE

<u>AP</u>		<u>AKORN INC</u>	<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>		<u>ASTRAL</u>	<u>EQ 250MG BASE/VIAL</u>	<u>A091049 001</u>	Jun 11, 2018
<u>AP</u>			<u>EQ 500MG BASE/VIAL</u>	<u>A091049 002</u>	Jun 11, 2018
<u>AP</u>			<u>EQ 1GM BASE/VIAL</u>	<u>A091049 003</u>	Jun 11, 2018
<u>AP</u>			<u>EQ 2GM BASE/VIAL</u>	<u>A091049 004</u>	Jun 11, 2018
<u>AP</u>		<u>HIKMA FARMACEUTICA</u>	<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>		<u>LUPIN</u>	<u>EQ 250MG BASE/VIAL</u>	<u>A065125 001</u>	Sep 30, 2003
<u>AP</u>			<u>EQ 500MG BASE/VIAL</u>	<u>A065125 002</u>	Sep 30, 2003
<u>AP</u>			<u>EQ 1GM BASE/VIAL</u>	<u>A065125 003</u>	Sep 30, 2003
<u>AP</u>			<u>EQ 2GM BASE/VIAL</u>	<u>A065125 004</u>	Sep 30, 2003
<u>AP</u>		<u>QILU</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>			<u>EQ 1GM BASE/VIAL</u>	<u>A203702 003</u>	Jun 29, 2016
<u>AP</u>			<u>EQ 2GM BASE/VIAL</u>	<u>A203702 004</u>	Jun 29, 2016
<u>AP</u>	<u>!</u>	<u>SANDOZ</u>	<u>EQ 250MG BASE/VIAL</u>	<u>A065169 001</u>	May 09, 2005
<u>AP</u>	<u>!</u>		<u>EQ 500MG BASE/VIAL</u>	<u>A065169 002</u>	May 09, 2005
<u>AP</u>	<u>!</u>		<u>EQ 1GM BASE/VIAL</u>	<u>A065169 003</u>	May 09, 2005
<u>AP</u>	<u>!</u>		<u>EQ 2GM BASE/VIAL</u>	<u>A065169 004</u>	May 09, 2005
<u>AP</u>		<u>WOCKHARDT</u>	<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>		<u>ALKEM LABS LTD</u>	<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>		<u>APOTEX</u>	<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>		<u>AUROBINDO PHARMA LTD</u>	<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>		<u>LUPIN</u>	<u>EQ 250MG BASE</u>	<u>A065135 001</u>	Jul 25, 2003
<u>AB</u>	<u>!</u>		<u>EQ 500MG BASE</u>	<u>A065135 002</u>	Jul 25, 2003
<u>AB</u>		<u>ORCHID HLTHCARE</u>	<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>		<u>WOCKHARDT</u>	<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

CEFUROXIME SODIUM

INJECTABLE; INJECTION

CEFUROXIME AND DEXTROSE IN DUPLEX CONTAINER

<u>AP</u>	<u>+</u> !	<u>B BRAUN</u>	<u>EQ 750MG BASE/VIAL</u>	<u>N050780 001</u>	Feb 21, 2001
<u>AP</u>	<u>+</u> !		<u>EQ 1.5GM BASE/VIAL</u>	<u>N050780 002</u>	Feb 21, 2001

CEFUROXIME SODIUM

<u>AP</u>		<u>ACS DOBFAR SPA</u>	<u>EQ 1.5GM BASE/VIAL</u>	<u>A064125 002</u>	May 30, 1997
<u>AP</u>			<u>EQ 7.5GM BASE/VIAL</u>	<u>A064124 001</u>	May 30, 1997
<u>AP</u>		<u>HIKMA FARMACEUTICA</u>	<u>EQ 1.5GM BASE/VIAL</u>	<u>A065048 002</u>	Jan 09, 2004
<u>AP</u>			<u>EQ 7.5GM BASE/VIAL</u>	<u>A065046 001</u>	Jan 09, 2004

PRESCRIPTION DRUG PRODUCT LIST

CEFUROXIME SODIUM

INJECTABLE; INJECTION

ZINACEF

<u>AP</u>	<u>+!</u>	<u>TELIGENT</u>	<u>EQ 1.5GM BASE/VIAL</u>	<u>N050558 003</u>	Oct 19, 1983
<u>AP</u>	<u>+!</u>		<u>EQ 7.5GM BASE/VIAL</u>	<u>N050558 004</u>	Oct 23, 1986

INJECTABLE; INTRAMUSCULAR, INTRAVENOUS

CEFUROXIME SODIUM

<u>AB</u>		<u>ACS DOBFAR SPA</u>	<u>EQ 750MG BASE/VIAL</u>	<u>A064125 001</u>	May 30, 1997
<u>AB</u>		<u>HIKMA FARMACEUTICA</u>	<u>EQ 750MG BASE/VIAL</u>	<u>A065048 001</u>	Jan 09, 2004

ZINACEF

<u>AB</u>	<u>+!</u>	<u>TELIGENT</u>	<u>EQ 750MG BASE/VIAL</u>	<u>N050558 002</u>	Oct 19, 1983
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CELECOXIB

CAPSULE; ORAL

CELEBREX

<u>AB</u>	<u>+</u>	<u>GD SEARLE</u>	<u>50MG</u>	<u>N020998 004</u>	Dec 15, 2006
<u>AB</u>	<u>+</u>		<u>100MG</u>	<u>N020998 001</u>	Dec 31, 1998
<u>AB</u>	<u>+</u>		<u>200MG</u>	<u>N020998 002</u>	Dec 31, 1998
<u>AB</u>	<u>+!</u>		<u>400MG</u>	<u>N020998 003</u>	Aug 29, 2002

CELECOXIB

<u>AB</u>		<u>ALEMBIC PHARMS LTD</u>	<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>		<u>AMNEAL PHARMS</u>	<u>50MG</u>	<u>A208833 001</u>	May 31, 2018
<u>AB</u>			<u>100MG</u>	<u>A208833 002</u>	May 31, 2018
<u>AB</u>			<u>200MG</u>	<u>A208833 003</u>	May 31, 2018
<u>AB</u>			<u>400MG</u>	<u>A208833 004</u>	May 31, 2018
<u>AB</u>		<u>APOTEX INC</u>	<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>		<u>AUROBINDO PHARMA LTD</u>	<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>		<u>CADILA PHARMS LTD</u>	<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>		<u>CIPLA</u>	<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>		<u>CSPC OUYI</u>	<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>		<u>JUBILANT GENERICS</u>	<u>50MG</u>	<u>A207061 001</u>	Apr 04, 2017
<u>AB</u>			<u>100MG</u>	<u>A207061 002</u>	Apr 04, 2017
<u>AB</u>			<u>200MG</u>	<u>A207061 003</u>	Apr 04, 2017
<u>AB</u>			<u>400MG</u>	<u>A207061 004</u>	Apr 04, 2017
<u>AB</u>		<u>LUPIN LTD</u>	<u>50MG</u>	<u>A202240 001</u>	Oct 29, 2014
<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>		<u>MACLEODS PHARMS LTD</u>	<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>		<u>MICRO LABS</u>	<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>		<u>MYLAN</u>	<u>50MG</u>	<u>A078857 001</u>	May 30, 2014
<u>AB</u>			<u>100MG</u>	<u>A078857 002</u>	Feb 11, 2015
<u>AB</u>			<u>200MG</u>	<u>A078857 003</u>	Feb 11, 2015
<u>AB</u>			<u>400MG</u>	<u>A078857 004</u>	Feb 11, 2015
<u>AB</u>		<u>QINGDAO BAHEAL PHARM</u>	<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>		<u>TEVA</u>	<u>50MG</u>	<u>A076898 001</u>	May 30, 2014

PRESCRIPTION DRUG PRODUCT LIST

CELECOXIB

CAPSULE; ORAL

CELECOXIB

<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 LABS PVT LTD	<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 PHARM LTD	<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

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>	AUROBINDO PHARMA LTD	<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>	BELCHER PHARMS	<u>EQ 250MG BASE</u>	<u>A062713 001</u>	Jul 15, 1988
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>A062713 002</u>	Jul 15, 1988
<u>AB</u>	HIKMA	<u>EQ 250MG BASE</u>	<u>A065215 001</u>	Jan 24, 2006
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>A065215 002</u>	Jan 24, 2006
<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>	ORCHID HLTHCARE	<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>	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
<u>AB</u>	YUNG SHIN PHARM	<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

KEFLEX

<u>AB</u>	+ PRAGMA	<u>EQ 250MG BASE</u>	<u>N050405 002</u>	
<u>AB</u>	+ PRAGMA	<u>EQ 500MG BASE</u>	<u>N050405 003</u>	
<u>AB</u>	+! PRAGMA	<u>EQ 750MG BASE</u>	<u>N050405 005</u>	May 12, 2006

CEPHALEXIN

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>	ORCHID HLTHCARE	<u>EQ 125MG BASE/5ML</u>	<u>A065326 001</u>	Jul 10, 2006
<u>AB</u>		<u>EQ 250MG BASE/5ML</u>	<u>A065326 002</u>	Jul 10, 2006
<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

TEVA
!

EQ 250MG BASE	A063023 001	Jan 12, 1989
EQ 500MG BASE	A063024 001	Jan 12, 1989

PRESCRIPTION DRUG PRODUCT LIST

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/DROPS; OPHTHALMIC

ZERVIAE

+! EYEVANCE EQ 0.24% BASE N208694 001 May 30, 2017

SYRUP; ORAL

CETIRIZINE HYDROCHLORIDE

AA	AMNEAL PHARMS	5MG/5ML	A090766 001	Oct 07, 2009
AA	BRECKENRIDGE	5MG/5ML	A078488 001	Oct 06, 2008
AA	LANNETT CO INC	5MG/5ML	A078876 001	May 11, 2012
AA	MLV	5MG/5ML	A090191 001	Nov 12, 2009
AA	! PERRIGO PHARMS CO	5MG/5ML	A078398 001	Jun 17, 2008
AA	TARO	5MG/5ML	A076601 001	Jun 20, 2008
AA	TEVA PHARMS	5MG/5ML	A077279 001	May 27, 2008

CETRORELIX

INJECTABLE; INJECTION

CETROTIDE

+! EMD SERONO INC EQ 0.25MG BASE/ML N021197 001 Aug 11, 2000

CEVIMELINE HYDROCHLORIDE

CAPSULE; ORAL

CEVIMELINE HYDROCHLORIDE

AB	HIKMA	30MG	A091591 001	Jul 08, 2013
AB	NOVEL LABS INC	30MG	A204746 001	Dec 30, 2016
AB	RISING	30MG	A203775 001	Jun 04, 2014

EVOXAC

AB	! DAIICHI SANKYO INC	30MG	N020989 002	Jan 11, 2000
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CHENODIOL

TABLET; ORAL

CHENODIOL

! NEXGEN PHARMA 250MG A091019 001 Oct 22, 2009

CHLORAMBUCIL

TABLET; ORAL

LEUKERAN

+! ASPEN GLOBAL INC 2MG N010669 002

CHLORAMPHENICOL SODIUM SUCCINATE

INJECTABLE; INJECTION

CHLORAMPHENICOL SODIUM SUCCINATE

! FRESENIUS KABI USA EQ 1GM BASE/VIAL A062365 001 Aug 25, 1982

CHLORDIAZEPOXIDE HYDROCHLORIDE

CAPSULE; ORAL

CHLORDIAZEPOXIDE HYDROCHLORIDE

AB	BARR	5MG	A084768 001	
AB		10MG	A083116 001	
AB		25MG	A084769 001	

LIBRIUM

AB	VALEANT PHARM INTL	5MG	A085461 001	
AB		10MG	A085472 001	
AB	!	25MG	A085475 001	

CHLORDIAZEPOXIDE HYDROCHLORIDE; CLIDINIUM BROMIDE

CAPSULE; ORAL

LIBRAX

+! VALEANT PHARMS 5MG; 2.5MG N012750 001

CHLORHEXIDINE GLUCONATE

SOLUTION; DENTAL

CHLORHEXIDINE GLUCONATE

AT	HI TECH PHARMA	0.12%	A074356 001	May 07, 1996
AT	LYNE	0.12%	A074291 001	Dec 28, 1995
AT	PHARM ASSOC	0.12%	A074522 001	Dec 15, 1995
AT	WOCKHARDT BIO AG	0.12%	A075006 001	Mar 03, 2004
AT	XTTRIUM	0.12%	A077789 001	Jun 18, 2009

PRESCRIPTION DRUG PRODUCT LISTCHLORHEXIDINE GLUCONATE

SOLUTION;DENTAL

PAROEX**AT** SUNSTAR AMERICAS **0.12%** **A076434 001** Nov 29, 2005PERIDEX**AT** +! 3M **0.12%** **N019028 001** Aug 13, 1986PERIOGARD**AT** COLGATE PALMOLIVE **0.12%** **A073695 001** Jan 14, 1994
CO**AT** COLGATE-PALMOLIVE **0.12%** **A203212 001** Jan 28, 2016
CO

TABLET;DENTAL

PERIOCHIP+! DEXCEL PHARMA 2.5MG **N020774 001** May 15, 1998CHLOROPROCAINE HYDROCHLORIDE

INJECTABLE;INJECTION

CHLOROPROCAINE HYDROCHLORIDE**AP** HOSPIRA **2%** **A087447 001** Apr 16, 1982**AP** **3%** **A087446 001** Apr 16, 1982**AP** WEST-WARD PHARMS **2%** **A040273 001** Sep 09, 1998
INT**AP** **3%** **A040273 002** Sep 09, 1998NESACAINE**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

NESACAINE

+! FRESENIUS KABI USA 1% **N009435 001**

SOLUTION;INTRATHECAL

CLOROTEKAL+ B BRAUN MEDICAL INC 50MG/5ML (10MG/ML) **N208791 001** Sep 26, 2017CHLOROQUINE PHOSPHATE

TABLET;ORAL

CHLOROQUINE PHOSPHATE**AA** HIKMA PHARMS **EQ 300MG BASE** **A083082 002** Sep 17, 1999**AA** IPCA LABS LTD **EQ 150MG BASE** **A090610 001** Dec 03, 2009**AA** **EQ 300MG BASE** **A090249 001** Dec 03, 2009**AA** ! NATCO PHARMA LTD **EQ 150MG BASE** **A091621 001** Jan 21, 2011**AA** ! **EQ 300MG BASE** **A090612 001** Jan 21, 2011CHLOROTHIAZIDE

SUSPENSION;ORAL

DIURIL+! SALIX PHARMS 250MG/5ML **N011870 001**

TABLET;ORAL

CHLOROTHIAZIDE! MYLAN 250MG **A084217 002**500MG **A084217 001**CHLOROTHIAZIDE SODIUM

INJECTABLE;INJECTION

CHLOROTHIAZIDE SODIUM**AP** AM REGENT **EQ 500MG BASE/VIAL** **A202561 001** Apr 22, 2013**AP** FRESENIUS KABI USA **EQ 500MG BASE/VIAL** **A090896 001** Oct 16, 2009**AP** MYLAN INSTITUTIONAL **EQ 500MG BASE/VIAL** **A202493 001** Jun 18, 2014**AP** SAGENT PHARMS INC **EQ 500MG BASE/VIAL** **A202462 001** May 29, 2015**AP** SUN PHARM **EQ 500MG BASE/VIAL** **A091546 001** Jul 26, 2011DIURIL**AP** +! OAK PHARMS AKORN **EQ 500MG BASE/VIAL** **N011145 005**CHLORPHENIRAMINE MALEATE; CODEINE PHOSPHATE

TABLET, EXTENDED RELEASE;ORAL

TUXARIN ERMAINPOINTE 8MG;54.3MG **N206323 001** Jun 22, 2015CHLORPHENIRAMINE MALEATE; HYDROCODONE BITARTRATE; PSEUDOEPHEDRINE HYDROCHLORIDE

SOLUTION;ORAL

HYDROCODONE BITARTRATE, CHLORPHENIRAMINE MALEATE AND PSEUDOEPHEDRINE HYDROCHLORIDE

! PADDOCK LLC 4MG/5ML;5MG/5ML;60MG/5ML **A204627 001** Apr 29, 2014

PRESCRIPTION DRUG PRODUCT LIST

CHLORPHENIRAMINE POLISTIREX; CODEINE POLISTIREX

SUSPENSION, EXTENDED RELEASE;ORAL

TUZISTRA XR

+! AYTU 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

AB TRIS PHARMA INC EQ 8MG MALEATE/5ML;EQ 10MG BITARTRATE/5ML **A091632 001** Oct 01, 2010

HYDROCODONE POLISTIREX AND CHLORPHENIRAMNE POLISTIREX

AB ! NEOS THERAP INC EQ 8MG MALEATE/5ML;EQ 10MG BITARTRATE/5ML **A091671 001** Jun 29, 2012

CHLORPROMAZINE HYDROCHLORIDE

INJECTABLE; INJECTION

CHLORPROMAZINE HYDROCHLORIDE

! WEST-WARD PHARMS 25MG/ML A083329 001
INT

TABLET;ORAL

CHLORPROMAZINE HYDROCHLORIDE

AB AMNEAL PHARMS CO 10MG **A209755 001** Sep 10, 2018

AB 25MG **A209755 002** Sep 10, 2018

AB 50MG **A209755 003** Sep 10, 2018

AB 100MG **A209755 004** Sep 10, 2018

AB 200MG **A209755 005** Sep 10, 2018

AB UPSHER SMITH LABS 10MG **A083386 001**

AB ! 25MG **A084112 001**

AB 50MG **A084113 001**

AB ! 100MG **A084114 001**

AB 200MG **A084115 001**

AB ZYDUS 10MG **A213368 001** Jan 17, 2020

AB 25MG **A213368 002** Jan 17, 2020

AB 50MG **A213368 003** Jan 17, 2020

AB 100MG **A213368 004** Jan 17, 2020

AB 200MG **A213368 005** Jan 17, 2020

CHLORTHALIDONE

TABLET;ORAL

CHLORTHALIDONE

AB ALKEM LABS LTD 25MG **A213412 001** Feb 11, 2020

AB 50MG **A213412 002** Feb 11, 2020

AB AMNEAL PHARMS CO 25MG **A207204 001** Jul 01, 2019

AB 50MG **A207204 002** Jul 01, 2019

AB ARISE 25MG **A210742 001** Oct 12, 2018

AB 50MG **A210742 002** Oct 12, 2018

AB MYLAN 25MG **A086831 002**

AB ! 50MG **A086831 001**

AB RICONPHARMA LLC 25MG **A206904 001** Mar 30, 2017

AB 50MG **A206904 002** Mar 30, 2017

AB SUN PHARM 25MG **A089286 002** Jul 21, 1986

INDUSTRIES

AB 50MG **A089286 001** Jul 21, 1986

AB UMEDICA LABS PVT 25MG **A207222 001** May 24, 2018

LTD

AB 50MG **A207222 002** May 24, 2018

AB UNICHEM LABS LTD 25MG **A211627 001** Aug 06, 2019

AB 50MG **A211627 002** Aug 06, 2019

AB VISTAPHARM 25MG **A211063 001** Feb 26, 2019

AB 50MG **A211063 002** Feb 26, 2019

AB ZYDUS PHARMS 25MG **A207813 001** May 10, 2019

AB 50MG **A207813 002** May 10, 2019

CHLORZOXAZONE

TABLET;ORAL

CHLORZOXAZONE

AA NOVITIUM PHARMA 500MG **A212254 001** Sep 12, 2019

AA ! WATSON LABS 500MG **A089859 001** May 04, 1988

AB ! MIKART 375MG **A040861 001** Jun 01, 2010

AB ! 750MG **A040861 002** Jun 01, 2010

AB NOVITIUM PHARMA 375MG **A212253 001** Nov 27, 2019

PRESCRIPTION DRUG PRODUCT LIST

3-93 (of 453)

CHLORZOXAZONE

TABLET; ORAL

CHLORZOXAZONE

<u>AB</u>		<u>750MG</u>	<u>A212253 002</u>	Nov 27, 2019
	!	MIKART	A207483 001	Jun 24, 2016

CHOLESTYRAMINE

POWDER; ORAL

CHOLESTYRAMINE

<u>AB</u>		ANI PHARMS INC	<u>EQ 4GM RESIN/PACKET</u>	<u>A074554 001</u>	Oct 02, 1996
<u>AB</u>			<u>EQ 4GM RESIN/SCOOPFUL</u>	<u>A074554 002</u>	Oct 02, 1996
<u>AB</u>		PAR PHARM	<u>EQ 4GM RESIN/PACKET</u>	<u>A077204 001</u>	Aug 26, 2005
<u>AB</u>			<u>EQ 4GM RESIN/SCOOPFUL</u>	<u>A077204 002</u>	Aug 26, 2005
<u>AB</u>	!	SANDOZ	<u>EQ 4GM RESIN/PACKET</u>	<u>A074557 001</u>	Aug 15, 1996
<u>AB</u>			<u>EQ 4GM RESIN/SCOOPFUL</u>	<u>A074557 002</u>	Aug 15, 1996
<u>AB</u>		ZYDUS PHARMS	<u>EQ 4GM RESIN/SCOOPFUL</u>	<u>A202901 001</u>	Jul 02, 2018

CHOLESTYRAMINE LIGHT

<u>AB</u>		PAR PHARM	<u>EQ 4GM RESIN/PACKET</u>	<u>A077203 001</u>	Aug 26, 2005
<u>AB</u>			<u>EQ 4GM RESIN/SCOOPFUL</u>	<u>A077203 002</u>	Aug 26, 2005
<u>AB</u>	!	SANDOZ	<u>EQ 4GM RESIN/PACKET</u>	<u>A074558 001</u>	Aug 15, 1996
<u>AB</u>			<u>EQ 4GM RESIN/SCOOPFUL</u>	<u>A074558 002</u>	Aug 15, 1996
<u>AB</u>		ZYDUS PHARMS	<u>EQ 4GM RESIN/SCOOPFUL</u>	<u>A202902 001</u>	Apr 25, 2017

PREVALITE

<u>AB</u>		UPSHER SMITH LABS	<u>EQ 4GM RESIN/PACKET</u>	<u>A073263 001</u>	Feb 22, 1996
<u>AB</u>			<u>EQ 4GM RESIN/SCOOPFUL</u>	<u>A073263 002</u>	Oct 30, 1997

CHOLIC ACID

CAPSULE; ORAL

CHOLBAM

+	RTRX	50MG	N205750 001	Mar 17, 2015
+	!	250MG	N205750 002	Mar 17, 2015

CHOLINE C-11

INJECTABLE; INTRAVENOUS

CHOLINE C-11

<u>AP</u>		DECATUR	<u>4-33.1mCi/ML</u>	<u>A206319 001</u>	Nov 13, 2015
<u>AP</u>	+	MCPRF	<u>4-33.1mCi/ML</u>	<u>N203155 001</u>	Sep 12, 2012
<u>AP</u>		UCSF RODIOPHARM	<u>4-33.1mCi/ML</u>	<u>A208444 001</u>	Nov 20, 2017
<u>AP</u>		WA UNIV SCH MED	<u>4-33.1mCi/ML</u>	<u>A208413 001</u>	Jan 10, 2017
		UNIV TX MD ANDERSON	4-100mCi/ML	A205690 001	Oct 29, 2015

CHOLINE FENOFIBRATE

CAPSULE, DELAYED RELEASE; ORAL

FENOFIBRIC ACID

<u>AB</u>		ACTAVIS ELIZABETH	<u>EQ 45MG FENOFIBRIC ACID</u>	<u>A200920 001</u>	Oct 07, 2015
<u>AB</u>			<u>EQ 135MG FENOFIBRIC ACID</u>	<u>A200920 002</u>	Oct 07, 2015
<u>AB</u>		ALEMBIC PHARMS LTD	<u>EQ 45MG FENOFIBRIC ACID</u>	<u>A208705 001</u>	May 12, 2017
<u>AB</u>			<u>EQ 135MG FENOFIBRIC ACID</u>	<u>A208705 002</u>	May 12, 2017
<u>AB</u>		ANCHEN PHARMS	<u>EQ 45MG FENOFIBRIC ACID</u>	<u>A201573 002</u>	Jul 18, 2013
<u>AB</u>			<u>EQ 135MG FENOFIBRIC ACID</u>	<u>A201573 001</u>	Jul 18, 2013
<u>AB</u>		AUROBINDO PHARMA LTD	<u>EQ 45MG FENOFIBRIC ACID</u>	<u>A212598 001</u>	Jul 25, 2019
<u>AB</u>			<u>EQ 135MG FENOFIBRIC ACID</u>	<u>A212598 002</u>	Jul 25, 2019
<u>AB</u>		GRAVITI PHARMS	<u>EQ 45MG FENOFIBRIC ACID</u>	<u>A211626 001</u>	Jul 18, 2019
<u>AB</u>			<u>EQ 135MG FENOFIBRIC ACID</u>	<u>A211626 002</u>	Jul 18, 2019
<u>AB</u>		IMPAX LABS INC	<u>EQ 45MG FENOFIBRIC ACID</u>	<u>A200264 001</u>	Sep 07, 2016
<u>AB</u>			<u>EQ 135MG FENOFIBRIC ACID</u>	<u>A200264 002</u>	Sep 07, 2016
<u>AB</u>		LUPIN LTD	<u>EQ 45MG FENOFIBRIC ACID</u>	<u>A200750 001</u>	Dec 04, 2013
<u>AB</u>			<u>EQ 135MG FENOFIBRIC ACID</u>	<u>A200750 002</u>	Dec 04, 2013

TRILIPIX

<u>AB</u>	+	ABBVIE	<u>EQ 45MG FENOFIBRIC ACID</u>	<u>N022224 001</u>	Dec 15, 2008
<u>AB</u>	+	!	<u>EQ 135MG FENOFIBRIC ACID</u>	<u>N022224 002</u>	Dec 15, 2008

CHORIOGONADOTROPIN ALFA

INJECTABLE; SUBCUTANEOUS

OVIDREL

+	!	EMD SERONO	EQ 0.25MG /0.5ML	N021149 002	Oct 06, 2003
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CHROMIC CHLORIDE

INJECTABLE; INJECTION

CHROMIC CHLORIDE IN PLASTIC CONTAINER

+	!	HOSPIRA	EQ 0.004MG CHROMIUM/ML	N018961 001	Jun 26, 1986
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PRESCRIPTION DRUG PRODUCT LISTCICLESONIDE

AEROSOL, METERED; INHALATION

ALVESCO

+	!	COVIS PHARMA BV	0.08MG/INH	N021658	002	Jan 10, 2008
+	!		0.16MG/INH	N021658	003	Jan 10, 2008

AEROSOL, METERED; NASAL

ZETONNA

+	!	COVIS PHARMA BV	0.037MG/INH	N202129	001	Jan 20, 2012
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SPRAY, METERED; NASAL

OMNARIS

+	!	COVIS PHARMA BV	0.05MG/INH	N022004	001	Oct 20, 2006
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CICLOPIROX

CREAM; TOPICAL

CICLOPIROX

AB		ACP NIMBLE	<u>0.77%</u>	A078463	001	Dec 20, 2010
AB		FOUGERA PHARMS	<u>0.77%</u>	A076435	001	Dec 29, 2004
AB		GLENMARK PHARMS	<u>0.77%</u>	A090273	001	Nov 10, 2009
AB		PERRIGO ISRAEL	<u>0.77%</u>	A077364	001	Mar 03, 2006
AB		TARO	<u>0.77%</u>	A076790	001	Apr 12, 2005

LOPROX

AB	+	!	MEDIMETRIKS PHARMS	<u>0.77%</u>	N018748	001	Dec 30, 1982
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GEL; TOPICAL

CICLOPIROX

AB	+	!	CNTY LINE PHARMS	<u>0.77%</u>	N020519	001	Jul 21, 1997
AB			FOUGERA PHARMS	<u>0.77%</u>	A077896	001	Jun 10, 2008
AB			GLENMARK GENERICS	<u>0.77%</u>	A091595	001	Feb 29, 2012
AB			PADDOCK LLC	<u>0.77%</u>	A078266	001	Jan 07, 2009

SHAMPOO; TOPICAL

CICLOPIROX

AT		ACTAVIS MID	<u>1%</u>	A090490	001	Nov 24, 2009
		ATLANTIC				
AT		FOUGERA PHARMS	<u>1%</u>	A090146	001	May 25, 2010
AT		PERRIGO CO	<u>1%</u>	A078594	001	Feb 16, 2010
AT		TARO	<u>1%</u>	A090269	001	Feb 23, 2011
AT		TELIGENT PHARMA INC	<u>1%</u>	A209975	001	Apr 05, 2018

LOPROX

AT	+	!	BAUSCH	<u>1%</u>	N021159	001	Feb 28, 2003
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SOLUTION; TOPICAL

CICLOPIROX

AT		ACELLA	<u>8%</u>	A078172	001	Sep 18, 2007
AT		ACP NIMBLE	<u>8%</u>	A078233	001	Sep 18, 2007
AT		ACTAVIS MID	<u>8%</u>	A078046	001	Sep 18, 2007
		ATLANTIC				
AT		AKORN	<u>8%</u>	A078975	001	Feb 17, 2010
AT		ANDA REPOSITORY	<u>8%</u>	A077687	001	Sep 18, 2007
AT		HI TECH PHARMA	<u>8%</u>	A078270	001	Sep 18, 2007
AT		NOVAST LABS	<u>8%</u>	A078124	001	Sep 18, 2007
AT		PERRIGO NEW YORK	<u>8%</u>	A077623	001	Sep 18, 2007
AT		TARO PHARM INDS	<u>8%</u>	A078144	001	Sep 18, 2007

PENLAC

AT	+	!	VALEANT BERMUDA	<u>8%</u>	N021022	001	Dec 17, 1999
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SUSPENSION; TOPICAL

CICLOPIROX

AB		FOUGERA PHARMS	<u>0.77%</u>	A076422	001	Aug 06, 2004
AB		PERRIGO NEW YORK	<u>0.77%</u>	A077676	001	Dec 15, 2006
AB		TARO	<u>0.77%</u>	A077092	001	Aug 10, 2005

LOPROX

AB	+	!	MEDIMETRIKS PHARMS	<u>0.77%</u>	N019824	001	Dec 30, 1988
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CIDOFOVIR

INJECTABLE; INJECTION

CIDOFOVIR

AP		EMCURE PHARMS LTD	<u>EQ 75MG BASE/ML</u>	A202501	001	Jul 26, 2012
AP	!	MYLAN INSTITUTIONAL	<u>EQ 75MG BASE/ML</u>	A201276	001	Jun 27, 2012

CILASTATIN SODIUM; IMIPENEM

POWDER; INTRAVENOUS

IMIPENEM AND CILASTATIN

AP		ACS DOBFAR	<u>EQ 500MG BASE/VIAL; 500MG/VIAL</u>	A090577	002	Dec 21, 2011	
AP		HQ SPCLT PHARMA	<u>EQ 500MG BASE/VIAL; 500MG/VIAL</u>	A207594	001	Dec 12, 2019	
AP	+	!	MERCK	<u>EQ 500MG BASE/VIAL; 500MG/VIAL</u>	N050587	002	Nov 26, 1985

PRESCRIPTION DRUG PRODUCT LISTCILASTATIN SODIUM; IMPENEM

POWDER; INTRAVENOUS

IMPENEM AND CILASTATIN

! ACS DOBFAR

EQ 250MG BASE/VIAL; 250MG/VIAL

A090577 001 Dec 21, 2011

CILASTATIN SODIUM; IMPENEM; RELEBACTAM

POWDER; INTRAVENOUS

RECARBRIO

+! MSD MERCK CO

EQ 500MG
BASE/VIAL; 500MG/VIAL; 250MG/VIAL

N212819 001 Jul 16, 2019

CILOSTAZOL

TABLET; ORAL

CILOSTAZOL

AB	APOTEX INC	50MG	A077030 001	Dec 10, 2004
AB		100MG	A077030 002	Dec 10, 2004
AB	BRECKENRIDGE PHARM	50MG	A077708 001	Sep 28, 2009
AB		100MG	A077708 002	Sep 28, 2009
AB	CASI PHARMS INC	50MG	A077310 001	Nov 08, 2005
AB		100MG	A077021 001	Nov 23, 2004
AB	CHARTWELL RX	50MG	A077722 001	Sep 24, 2012
AB		100MG	A077831 001	Sep 24, 2012
AB	HIKMA	50MG	A077024 001	May 17, 2005
AB		100MG	A077024 002	May 17, 2005
AB	SLATE	50MG	A077208 002	Mar 29, 2006
AB		100MG	A077208 001	Mar 29, 2006
AB	! TEVA	50MG	A077027 001	Nov 24, 2004
AB	!	100MG	A077027 002	Nov 24, 2004

CIMETIDINE

TABLET; ORAL

CIMETIDINE

AB	HIKMA	200MG	A074890 001	Dec 18, 1998
AB		300MG	A074890 002	Dec 18, 1998
AB		400MG	A074890 003	Dec 18, 1998
AB		800MG	A074890 004	Dec 18, 1998
AB	MYLAN	200MG	A074246 001	May 17, 1994
AB		300MG	A074246 002	May 17, 1994
AB		400MG	A074246 003	May 17, 1994
AB	!	800MG	A074246 004	May 17, 1994
AB	PLIVA	800MG	A074566 001	Feb 27, 1997
AB	TEVA	200MG	A074151 001	May 17, 1994
AB		300MG	A074151 002	May 17, 1994
AB		400MG	A074151 003	May 17, 1994
AB		800MG	A074463 001	May 17, 1994

CIMETIDINE HYDROCHLORIDE

INJECTABLE; INJECTION

CIMETIDINE HYDROCHLORIDE

DAVA PHARMS INC

EQ 300MG BASE/2ML

A074428 001 Apr 25, 1996

SOLUTION; ORAL

CIMETIDINE HYDROCHLORIDE

AA	! HI TECH PHARMA	EQ 300MG BASE/5ML	A074664 001	Oct 28, 1997
AA	PHARM ASSOC	EQ 300MG BASE/5ML	A074553 001	Jan 27, 1997
AA	WOCKHARDT BIO AG	EQ 300MG BASE/5ML	A074757 001	Oct 17, 1997

CINACALCET HYDROCHLORIDE

TABLET; ORAL

CINACALCET HYDROCHLORIDE

AB	ALKEM LABS LTD	EQ 30MG BASE	A210570 001	May 17, 2019
AB		EQ 60MG BASE	A210570 002	May 17, 2019
AB		EQ 90MG BASE	A210570 003	May 17, 2019
AB	AUROBINDO PHARMA LTD	EQ 30MG BASE	A206125 001	Mar 08, 2018
AB		EQ 60MG BASE	A206125 002	Mar 08, 2018
AB		EQ 90MG BASE	A206125 003	Mar 08, 2018
AB	CIPLA	EQ 30MG BASE	A208915 001	Mar 08, 2018
AB		EQ 60MG BASE	A208915 002	Mar 08, 2018
AB		EQ 90MG BASE	A208915 003	Mar 08, 2018
AB	MYLAN	EQ 30MG BASE	A203422 001	Oct 16, 2018
AB		EQ 60MG BASE	A203422 002	Oct 16, 2018
AB		EQ 90MG BASE	A203422 003	Oct 16, 2018
AB	PIRAMAL HLTHCARE UK	EQ 30MG BASE	A210207 001	Aug 01, 2018
AB		EQ 60MG BASE	A210207 002	Aug 01, 2018
AB		EQ 90MG BASE	A210207 003	Aug 01, 2018

PRESCRIPTION DRUG PRODUCT LIST

CINACALCET HYDROCHLORIDE

TABLET; ORAL

CINACALCET HYDROCHLORIDE

<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 30MG BASE</u>	<u>A207008 001</u>	Oct 11, 2018
<u>AB</u>			<u>EQ 60MG BASE</u>	<u>A207008 002</u>	Oct 11, 2018
<u>AB</u>			<u>EQ 90MG BASE</u>	<u>A207008 003</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
<u>SENSIPAR</u>					
<u>AB</u>	+	AMGEN	<u>EQ 30MG BASE</u>	<u>N021688 001</u>	Mar 08, 2004
<u>AB</u>	+		<u>EQ 60MG BASE</u>	<u>N021688 002</u>	Mar 08, 2004
<u>AB</u>	+		<u>EQ 90MG BASE</u>	<u>N021688 003</u>	Mar 08, 2004

CIPROFLOXACIN

FOR SUSPENSION; ORAL

CIPRO

+	BAYER HLTHCARE	250MG/5ML	N020780 001	Sep 26, 1997
+	!	500MG/5ML	N020780 002	Sep 26, 1997

INJECTABLE; INJECTION

CIPROFLOXACIN

<u>AP</u>	!	BAXTER HLTHCARE CORP	<u>200MG/20ML (10MG/ML)</u>	<u>A078062 001</u>	Apr 29, 2008
<u>AP</u>	!		<u>400MG/40ML (10MG/ML)</u>	<u>A078062 002</u>	Apr 29, 2008
<u>AP</u>		HIKMA FARMACEUTICA	<u>200MG/20ML (10MG/ML)</u>	<u>A076717 001</u>	Dec 22, 2009
<u>AP</u>			<u>400MG/40ML (10MG/ML)</u>	<u>A076717 002</u>	Dec 22, 2009
<u>AP</u>		HOSPIRA	<u>200MG/20ML (10MG/ML)</u>	<u>A077245 001</u>	Aug 28, 2006
<u>AP</u>			<u>400MG/40ML (10MG/ML)</u>	<u>A077245 002</u>	Aug 28, 2006

CIPROFLOXACIN IN DEXTROSE 5% IN PLASTIC CONTAINER

<u>AP</u>		BAXTER HLTHCARE CORP	<u>200MG/100ML</u>	<u>A078024 001</u>	Mar 18, 2008
<u>AP</u>			<u>400MG/200ML</u>	<u>A078024 002</u>	Mar 18, 2008
<u>AP</u>		HIKMA FARMACEUTICA	<u>400MG/200ML</u>	<u>A078431 001</u>	Nov 18, 2009
<u>AP</u>	!	HOSPIRA	<u>200MG/100ML</u>	<u>A077753 001</u>	Mar 18, 2008
<u>AP</u>	!		<u>400MG/200ML</u>	<u>A077753 002</u>	Mar 18, 2008
<u>AP</u>		INFORLIFE	<u>200MG/100ML</u>	<u>A078252 001</u>	Mar 18, 2008
<u>AP</u>			<u>400MG/200ML</u>	<u>A078252 002</u>	Mar 18, 2008

INJECTABLE, SUSPENSION; OTIC

OTIPRIO

+	!	OTONOMY INC	6% (60MG/ML)	N207986 001	Dec 10, 2015
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CIPROFLOXACIN HYDROCHLORIDE

OINTMENT; OPHTHALMIC

CILOXAN

+	!	NOVARTIS	EQ 0.3% BASE	N020369 001	Mar 30, 1998
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SOLUTION/DROPS; OPHTHALMIC

CILOXAN

<u>AT</u>	+	!	NOVARTIS	<u>EQ 0.3% BASE</u>	<u>N019992 001</u>	Dec 31, 1990
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CIPROFLOXACIN HYDROCHLORIDE

<u>AT</u>		AKORN INC	<u>EQ 0.3% BASE</u>	<u>A076555 001</u>	Dec 11, 2008
<u>AT</u>		ALTAIRE PHARMS INC	<u>EQ 0.3% BASE</u>	<u>A204613 001</u>	May 03, 2018
<u>AT</u>		FDC LTD	<u>EQ 0.3% BASE</u>	<u>A077568 001</u>	Jun 30, 2008
<u>AT</u>		RISING	<u>EQ 0.3% BASE</u>	<u>A077689 001</u>	Dec 13, 2006
<u>AT</u>		TELGENT	<u>EQ 0.3% BASE</u>	<u>A076754 001</u>	Jun 09, 2004
<u>AT</u>		WATSON LABS INC	<u>EQ 0.3% BASE</u>	<u>A076673 001</u>	Jan 21, 2005

SOLUTION/DROPS; OTIC

CETRAXAL

+	!	WRASER PHARMS	EQ 0.2% BASE	N021918 001	May 01, 2009
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TABLET; ORAL

CIPRO

<u>AB</u>	+	BAYER HLTHCARE	<u>EQ 250MG BASE</u>	<u>N019537 002</u>	Oct 22, 1987
<u>AB</u>	+	!	<u>EQ 500MG BASE</u>	<u>N019537 003</u>	Oct 22, 1987

CIPROFLOXACIN HYDROCHLORIDE

<u>AB</u>		AUROBINDO PHARMA	<u>EQ 250MG BASE</u>	<u>A077859 001</u>	Apr 26, 2007
<u>AB</u>			<u>EQ 500MG BASE</u>	<u>A077859 002</u>	Apr 26, 2007
<u>AB</u>			<u>EQ 750MG BASE</u>	<u>A077859 003</u>	Apr 26, 2007
<u>AB</u>		CARLSBAD	<u>EQ 250MG BASE</u>	<u>A076126 002</u>	Jun 09, 2004
<u>AB</u>			<u>EQ 500MG BASE</u>	<u>A076126 003</u>	Jun 09, 2004
<u>AB</u>			<u>EQ 750MG BASE</u>	<u>A076126 004</u>	Jun 09, 2004
<u>AB</u>		CHARTWELL	<u>EQ 250MG BASE</u>	<u>A076896 001</u>	Nov 04, 2004
<u>AB</u>			<u>EQ 500MG BASE</u>	<u>A076896 002</u>	Nov 04, 2004

PRESCRIPTION DRUG PRODUCT LIST

3-97 (of 453)

CIPROFLOXACIN HYDROCHLORIDE

TABLET; ORAL

CIPROFLOXACIN HYDROCHLORIDE

<u>AB</u>		<u>EQ 750MG BASE</u>	<u>A076896 003</u>	Nov 04, 2004
<u>AB</u>	DR REDDYS LABS LTD	<u>EQ 100MG BASE</u>	<u>A075593 002</u>	Jun 09, 2004
<u>AB</u>		<u>EQ 250MG BASE</u>	<u>A075593 003</u>	Jun 09, 2004
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>A075593 004</u>	Jun 09, 2004
<u>AB</u>		<u>EQ 750MG BASE</u>	<u>A075593 001</u>	Jun 09, 2004
<u>AB</u>	HIKMA	<u>EQ 250MG BASE</u>	<u>A076558 002</u>	Jun 09, 2004
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>A076558 003</u>	Jun 09, 2004
<u>AB</u>		<u>EQ 750MG BASE</u>	<u>A076558 004</u>	Jun 09, 2004
<u>AB</u>	IVAX SUB TEVA PHARMS	<u>EQ 250MG BASE</u>	<u>A076089 002</u>	Jun 09, 2004
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>A076089 003</u>	Jun 09, 2004
<u>AB</u>		<u>EQ 750MG BASE</u>	<u>A076089 004</u>	Jun 09, 2004
<u>AB</u>	MYLAN	<u>EQ 500MG BASE</u>	<u>A075817 003</u>	Jun 09, 2004
<u>AB</u>	TARO	<u>EQ 100MG BASE</u>	<u>A076912 001</u>	Feb 18, 2005
<u>AB</u>		<u>EQ 250MG BASE</u>	<u>A076912 002</u>	Oct 06, 2004
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>A076912 003</u>	Oct 06, 2004
<u>AB</u>		<u>EQ 750MG BASE</u>	<u>A076912 004</u>	Oct 06, 2004
<u>AB</u>	UNIQUE PHARM LABS	<u>EQ 250MG BASE</u>	<u>A076639 001</u>	Sep 10, 2004
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>A076639 002</u>	Sep 10, 2004
<u>AB</u>		<u>EQ 750MG BASE</u>	<u>A076639 003</u>	Sep 10, 2004
<u>AB</u>	WATSON LABS	<u>EQ 100MG BASE</u>	<u>A076794 001</u>	Feb 10, 2005
<u>AB</u>		<u>EQ 250MG BASE</u>	<u>A076794 002</u>	Jun 09, 2004
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>A076794 003</u>	Jun 09, 2004
<u>AB</u>		<u>EQ 750MG BASE</u>	<u>A076794 004</u>	Jun 09, 2004
<u>AB</u>	YILING PHARM LTD	<u>EQ 250MG BASE</u>	<u>A208921 001</u>	Jun 22, 2018
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>A208921 002</u>	Jun 22, 2018

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

+! NOVARTIS EQ 0.2% BASE;1% N020805 001 Feb 10, 1998

CIPROFLOXACIN; CIPROFLOXACIN HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL

CIPROFLOXACIN EXTENDED RELEASE

<u>AB</u>	!	ANCHEN PHARMS	<u>425.2MG;EQ 574.9MG BASE</u>	<u>A078166 001</u>	Nov 27, 2007
<u>AB</u>		DR REDDYS LABS LTD	<u>425.2MG;EQ 574.9MG BASE</u>	<u>A077701 001</u>	Mar 26, 2007
	!	ANCHEN PHARMS	212.6MG;EQ 287.5MG BASE	A078166 002	Nov 27, 2007

CIPROFLOXACIN; DEXAMETHASONE

SUSPENSION/DROPS; OTIC

CIPRODEX

+! NOVARTIS 0.3%;0.1% N021537 001 Jul 18, 2003

CISATRACURIUM BESYLATE

INJECTABLE; INJECTION

CISATRACURIUM BESYLATE

<u>AP</u>		FRESENIUS KABI USA	<u>EQ 2MG BASE/ML</u>	<u>A203183 001</u>	Feb 26, 2015
<u>AP</u>		HOSPIRA INC	<u>EQ 2MG BASE/ML</u>	<u>A203236 001</u>	Mar 30, 2018
<u>AP</u>			<u>EQ 2MG BASE/ML</u>	<u>A203238 001</u>	Mar 30, 2018
<u>AP</u>			<u>EQ 10MG BASE/ML</u>	<u>A203236 002</u>	Mar 30, 2018
<u>AP</u>		JIANGSU HENGRUI MED	<u>EQ 2MG BASE/ML</u>	<u>A209334 001</u>	Aug 30, 2017
<u>AP</u>		MEITHEAL	<u>EQ 2MG BASE/ML</u>	<u>A211668 001</u>	Apr 25, 2019
<u>AP</u>			<u>EQ 2MG BASE/ML</u>	<u>A211669 001</u>	Apr 25, 2019
<u>AP</u>			<u>EQ 10MG BASE/ML</u>	<u>A211668 002</u>	Apr 25, 2019
<u>AP</u>		SANDOZ INC	<u>EQ 2MG BASE/ML</u>	<u>A200159 001</u>	Feb 03, 2012
<u>AP</u>		SOMERSET	<u>EQ 2MG BASE/ML</u>	<u>A209132 001</u>	Apr 24, 2019
<u>AP</u>		ZYDUS PHARMS	<u>EQ 2MG BASE/ML</u>	<u>A212171 001</u>	Nov 04, 2019
<u>AP</u>			<u>EQ 10MG BASE/ML</u>	<u>A212171 002</u>	Nov 04, 2019

CISATRACURIUM BESYLATE PRESERVATIVE FREE

<u>AP</u>		FRESENIUS KABI USA	<u>EQ 2MG BASE/ML</u>	<u>A203182 001</u>	Feb 26, 2015
<u>AP</u>			<u>EQ 10MG BASE/ML</u>	<u>A203182 002</u>	Feb 26, 2015
<u>AP</u>		JIANGSU HENGRUI MED	<u>EQ 2MG BASE/ML</u>	<u>A204960 001</u>	Jan 27, 2017
<u>AP</u>			<u>EQ 10MG BASE/ML</u>	<u>A204960 002</u>	Sep 19, 2017
<u>AP</u>		SANDOZ INC	<u>EQ 2MG BASE/ML</u>	<u>A200154 001</u>	Feb 03, 2012
<u>AP</u>			<u>EQ 10MG BASE/ML</u>	<u>A200154 002</u>	Feb 03, 2012
<u>AP</u>		SOMERSET THERAPS	<u>EQ 2MG BASE/ML</u>	<u>A206791 001</u>	Feb 20, 2019

PRESCRIPTION DRUG PRODUCT LIST

CISATRACURIUM BESYLATE

INJECTABLE; INJECTION

CISATRACURIUM BESYLATE PRESERVATIVE FREE

LLC

<u>AP</u>		<u>EQ 10MG BASE/ML</u>	<u>A206791 002</u>	Feb 20, 2019
	<u>NIMBEX</u>			
<u>AP</u>	+! ABBVIE	<u>EQ 2MG BASE/ML</u>	<u>N020551 001</u>	Dec 15, 1995
	<u>NIMBEX PRESERVATIVE FREE</u>			
<u>AP</u>	+! ABBVIE	<u>EQ 2MG BASE/ML</u>	<u>N020551 003</u>	Dec 15, 1995
<u>AP</u>	+!	<u>EQ 10MG BASE/ML</u>	<u>N020551 002</u>	Dec 15, 1995

CISPLATIN

INJECTABLE; INJECTION

CISPLATIN

<u>AP</u>	ACCORD HLTHCARE	<u>1MG/ML</u>	<u>A206774 001</u>	Aug 18, 2015
<u>AP</u>	! FRESENIUS KABI USA	<u>1MG/ML</u>	<u>A074735 001</u>	Jul 16, 1999
<u>AP</u>	GLAND PHARMA LTD	<u>1MG/ML</u>	<u>A207323 001</u>	Mar 17, 2017
<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>	WEST-WARD PHARMS INT	<u>1MG/ML</u>	<u>A075036 001</u>	Nov 07, 2000

CITALOPRAM HYDROBROMIDE

SOLUTION; ORAL

CITALOPRAM HYDROBROMIDE

<u>AA</u>	AUROBINDO PHARMA LTD	<u>EQ 10MG BASE/5ML</u>	<u>A077812 001</u>	Aug 28, 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
<u>AA</u>	LANNETT CO INC	<u>EQ 10MG BASE/5ML</u>	<u>A077629 001</u>	Jun 15, 2006

TABLET; ORAL

CELEXA

<u>AB</u>	+ ALLERGAN	<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>	ACP NIMBLE	<u>EQ 10MG BASE</u>	<u>A077048 001</u>	Nov 16, 2004
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A077048 002</u>	Nov 16, 2004
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A077048 003</u>	Nov 16, 2004
<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 INC	<u>EQ 10MG BASE</u>	<u>A077046 001</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 GENERICS	<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>	JUBILANT GENERICS	<u>EQ 10MG BASE</u>	<u>A205407 001</u>	Dec 23, 2015
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A205407 002</u>	Dec 23, 2015
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A205407 003</u>	Dec 23, 2015
<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>	TORPHARM	<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>	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

PRESCRIPTION DRUG PRODUCT LISTCITRIC ACID; GLUCONOLACTONE; MAGNESIUM CARBONATE

SOLUTION; IRRIGATION

RENACIDIN

+! UNITED GUARDIAN 6.602GM/100ML; 198MG/100ML; 3.177GM/100ML N019481 001 Oct 02, 1990

CITRIC ACID; MAGNESIUM OXIDE; SODIUM PICOSULFATE

SOLUTION; ORAL

CLENPIQ

+! FERRING PHARMS INC 12GM/BOT; 3.5GM/BOT; 10MG/BOT N209589 001 Nov 28, 2017

CITRIC ACID; UREA C-13

FOR SOLUTION, TABLET, FOR SOLUTION; ORAL

IDKIT:HP

+! EXALENZ BIOSCIENCE N/A, 4GM; 75MG, N/A N021314 001 Dec 17, 2002

CLADRIBINE

INJECTABLE; INJECTION

CLADRIBINE**AP** ! FRESENIUS KABI USA 1MG/ML **A076571 001** Apr 22, 2004**AP** HISUN PHARM 1MG/ML **A210856 001** Nov 25, 2019

HANGZHOU

AP MYLAN LABS LTD 1MG/ML **A200510 001** Oct 06, 2011**AP** WEST-WARD PHARMS 1MG/ML **A075405 001** Feb 28, 2000

INT

TABLET; ORAL

MAVENCLAD

+! EMD SERONO INC 10MG N022561 001 Mar 29, 2019

CLARITHROMYCIN

FOR SUSPENSION; ORAL

CLARITHROMYCIN**AB** SANDOZ 125MG/5ML **A065283 002** Sep 04, 2007**AB** ! 250MG/5ML **A065283 003** Sep 04, 2007

TABLET; ORAL

CLARITHROMYCIN**AB** ! AUROBINDO 250MG **A065489 001** Jul 25, 2012**AB** ! 500MG **A065489 002** Jul 25, 2012**AB** CHARTWELL 250MG **A065384 001** Aug 20, 2007**AB** 500MG **A065384 002** Aug 20, 2007**AB** HEC PHARM 250MG **A203584 001** Sep 28, 2015**AB** 500MG **A203584 002** Sep 28, 2015**AB** HIKMA 250MG **A065178 002** May 25, 2004**AB** 500MG **A065178 001** May 25, 2004**AB** SANDOZ 250MG **A065144 001** Oct 18, 2005**AB** 500MG **A065136 001** Aug 25, 2005**AB** STRIDES PHARMA 250MG **A202710 001** Jun 10, 2013**AB** 500MG **A202710 002** Jun 10, 2013**AB** TEVA 250MG **A065155 001** May 31, 2005**AB** 500MG **A065155 002** May 31, 2005**AB** WOCKHARDT 250MG **A065266 001** May 31, 2006**AB** 500MG **A065266 002** May 31, 2006

TABLET, EXTENDED RELEASE; ORAL

CLARITHROMYCIN**AB** ACTAVIS LABS FL INC 500MG **A065145 001** Jun 24, 2004**AB** ! MAYNE PHARMA 500MG **A065154 001** May 18, 2005**AB** NOSTRUM LABS INC 500MG **A203243 001** Feb 29, 2016**AB** SUNSHINE LAKE 500MG **A208987 001** Jul 09, 2018CLEMASTINE FUMARATE

SYRUP; ORAL

CLEMASTINE FUMARATE

! TEVA

EQ 0.5MG BASE/5ML A073399 001 Jun 30, 1994

TABLET; ORAL

CLEMASTINE FUMARATE

! TEVA

2.68MG A073283 001 Jan 31, 1992

CLEVIDIPINE

EMULSION; INTRAVENOUS

CLEVIPREX

+! CHIESI USA INC 25MG/50ML (0.5MG/ML) N022156 001 Aug 01, 2008

+! 50MG/100ML (0.5MG/ML) N022156 002 Aug 01, 2008

PRESCRIPTION DRUG PRODUCT LIST

CLINDAMYCIN HYDROCHLORIDE

CAPSULE; ORAL

CLEOCIN HYDROCHLORIDE

AB	+	PHARMACIA AND UPJOHN	EQ 75MG BASE	N050162 001	
AB	+		EQ 150MG BASE	N050162 002	
AB	+	!	EQ 300MG BASE	N050162 003	Apr 14, 1988
<u>CLINDAMYCIN HYDROCHLORIDE</u>					
AB		ACP NIMBLE	EQ 150MG BASE	A063029 001	Sep 20, 1989
AB			EQ 300MG BASE	A063029 002	Aug 05, 2005
AB		AUROBINDO PHARMA	EQ 150MG BASE	A065442 001	Aug 26, 2009
AB			EQ 300MG BASE	A065442 002	Aug 26, 2009
AB		EPIC PHARMA LLC	EQ 150MG BASE	A065194 001	Mar 22, 2004
AB			EQ 300MG BASE	A065194 002	Mar 22, 2004
AB		LANNETT CO INC	EQ 75MG BASE	A065243 002	Aug 12, 2005
AB			EQ 150MG BASE	A065243 003	Aug 12, 2005
AB			EQ 300MG BASE	A065243 001	Aug 12, 2005
AB		MICRO LABS	EQ 75MG BASE	A207402 001	Nov 05, 2018
AB			EQ 150MG BASE	A207402 002	Nov 05, 2018
AB			EQ 300MG BASE	A207402 003	Nov 05, 2018
AB		SUN PHARM INDS LTD	EQ 150MG BASE	A065061 001	Feb 02, 2001
AB			EQ 300MG BASE	A065061 002	Feb 02, 2001
AB		WATSON LABS	EQ 150MG BASE	A063083 001	Jul 31, 1991
AB			EQ 300MG BASE	A063083 002	Mar 18, 2003
AB		ZYDUS PHARMS USA	EQ 75MG BASE	A065217 001	Jan 31, 2005
AB			EQ 150MG BASE	A065217 002	Jan 31, 2005
AB			EQ 300MG BASE	A065217 003	Jan 31, 2005

CLINDAMYCIN PALMITATE HYDROCHLORIDE

FOR SOLUTION; ORAL

CLEOCIN

AA	!	PHARMACIA AND UPJOHN	EQ 75MG BASE/5ML	A062644 001	Apr 07, 1986
<u>CLINDAMYCIN PALMITATE HYDROCHLORIDE</u>					
AA		AMNEAL PHARMS	EQ 75MG BASE/5ML	A203513 001	Mar 13, 2014
AA		AUROBINDO PHARMA LTD	EQ 75MG BASE/5ML	A202409 001	Apr 30, 2013
AA		HERITAGE PHARMS INC	EQ 75MG BASE/5ML	A207047 001	May 11, 2018
AA		LYNE	EQ 75MG BASE/5ML	A201821 001	Aug 28, 2012
AA		ORIT LABS LLC	EQ 75MG BASE/5ML	A206958 001	May 05, 2017
AA		PADDOCK LLC	EQ 75MG BASE/5ML	A090902 001	Jul 07, 2010

CLINDAMYCIN PHOSPHATE

AEROSOL, FOAM; TOPICAL

CLINDAMYCIN PHOSPHATE

AT		PERRIGO UK FINCO	1%	A090785 001	Mar 31, 2010
AT		TARO PHARM INDS LTD	1%	A210004 001	Mar 11, 2020
<u>EVOCLIN</u>					
AT	+	MYLAN	1%	N050801 001	Oct 22, 2004
CREAM; VAGINAL					
<u>CLEOCIN</u>					
AB	+	PHARMACIA AND UPJOHN	EQ 2% BASE	N050680 002	Mar 02, 1998
<u>CLINDAMYCIN PHOSPHATE</u>					
AB		FOUGERA PHARMS	EQ 2% BASE	A065139 001	Dec 27, 2004
CLINDESSE					
	+	PERRIGO PHARMA INTL	EQ 2% BASE	N050793 001	Nov 30, 2004
GEL; TOPICAL					
<u>CLEOCIN T</u>					
AB	+	PHARMACIA AND UPJOHN	EQ 1% BASE	N050615 001	Jan 07, 1987
<u>CLINDAMYCIN PHOSPHATE</u>					
AB		FOUGERA PHARMS	EQ 1% BASE	A064160 001	Jan 28, 2000
CLINDAGEL					
BT	+	BAUSCH	EQ 1% BASE	N050782 001	Nov 27, 2000
INJECTABLE; INJECTION					
<u>CLEOCIN PHOSPHATE</u>					
AP		PHARMACIA AND UPJOHN	EQ 150MG BASE/ML	A062803 001	Oct 16, 1987
AP	+	!	EQ 150MG BASE/ML	N050441 001	
<u>CLEOCIN PHOSPHATE IN DEXTROSE 5% IN PLASTIC CONTAINER</u>					
AP	+	PHARMACIA AND UPJOHN	EQ 6MG BASE/ML	N050639 001	Aug 30, 1989
AP	+		EQ 12MG BASE/ML	N050639 002	Aug 30, 1989
AP	+		EQ 18MG BASE/ML	N050639 003	Apr 10, 1991

PRESCRIPTION DRUG PRODUCT LIST

CLINDAMYCIN PHOSPHATE

INJECTABLE; INJECTION

CLINDAMYCIN PHOSPHATE

<u>AP</u>	ALVOGEN INC	<u>EQ 150MG BASE/ML</u>	<u>A062800 001</u>	Jul 24, 1987
<u>AP</u>		<u>EQ 150MG BASE/ML</u>	<u>A062801 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>	MYLAN LABS LTD	<u>EQ 150MG BASE/ML</u>	<u>A204748 001</u>	Oct 10, 2017
<u>AP</u>		<u>EQ 150MG BASE/ML</u>	<u>A204749 001</u>	Oct 10, 2017
<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
<u>AP</u>	WEST-WARD PHARMS INT	<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

CLINDAMYCIN PHOSPHATE IN 5% DEXTROSE IN PLASTIC CONTAINER

<u>AP</u>	AKORN INC	<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>	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>	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%

+	ABRAXIS PHARM	EQ 900MG BASE/100ML	N050635 001	Dec 22, 1989
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LOTION; TOPICAL

CLEOCIN T

<u>AB</u>	+	PHARMACIA AND UPJOHN	<u>EQ 1% BASE</u>	<u>N050600 001</u>	May 31, 1989
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CLINDAMYCIN PHOSPHATE

<u>AB</u>	FOUGERA PHARMS	<u>EQ 1% BASE</u>	<u>A065067 001</u>	Jan 31, 2002
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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

CLEOCIN T

<u>AT</u>	+	PHARMACIA AND UPJOHN	<u>EQ 1% BASE</u>	<u>N050537 001</u>
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CLINDA-DERM

<u>AT</u>	PADDOCK LLC	<u>EQ 1% BASE</u>	<u>A063329 001</u>	Sep 30, 1992
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CLINDAMYCIN PHOSPHATE

<u>AT</u>	CADILA	<u>EQ 1% BASE</u>	<u>A208767 001</u>	Jul 16, 2018
<u>AT</u>	ENCUBE ETHICALS	<u>EQ 1% BASE</u>	<u>A209914 001</u>	Jan 28, 2019
<u>AT</u>	FOUGERA PHARMS	<u>EQ 1% BASE</u>	<u>A065254 001</u>	Feb 14, 2006
<u>AT</u>	FOUGERA PHARMS INC	<u>EQ 1% BASE</u>	<u>A064159 001</u>	Jun 05, 1997
<u>AT</u>	G AND W LABS INC	<u>EQ 1% BASE</u>	<u>A062811 001</u>	Sep 01, 1988
<u>AT</u>	GLASSHOUSE PHARMS	<u>EQ 1% BASE</u>	<u>A209846 001</u>	Feb 08, 2018
<u>AT</u>	PERRIGO NEW YORK	<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>	TELIGENT PHARMA INC	<u>EQ 1% BASE</u>	<u>A206945 001</u>	Dec 30, 2016
<u>AT</u>	VINTAGE PHARMS	<u>EQ 1% BASE</u>	<u>A203343 001</u>	May 29, 2015

SUPPOSITORY; VAGINAL

CLEOCIN

+	PHARMACIA AND UPJOHN	100MG	N050767 001	Aug 13, 1999
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SWAB; TOPICAL

CLEOCIN

<u>AT</u>	+	PHARMACIA AND UPJOHN	<u>EQ 1% BASE</u>	<u>N050537 002</u>	Feb 22, 1994
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CLINDAMYCIN PHOSPHATE

<u>AT</u>	AKORN	<u>EQ 1% BASE</u>	<u>A065513 001</u>	Jun 17, 2010
<u>AT</u>	PERRIGO NEW YORK	<u>EQ 1% BASE</u>	<u>A065049 001</u>	May 25, 2000

CLINDETS

<u>AT</u>	PERRIGO CO	<u>EQ 1% BASE</u>	<u>A064136 001</u>	Sep 30, 1996
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PRESCRIPTION DRUG PRODUCT LIST

CLINDAMYCIN PHOSPHATE; TRETINOIN

GEL; TOPICAL

CLINDAMYCIN PHOSPHATE AND TRETINOIN

AB	ACTAVIS MID ATLANTIC	1.2%;0.025%	A202564 001	Jun 12, 2015
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ZIANA

AB	+! MEDICIS	1.2%;0.025%	N050802 001	Nov 07, 2006
	VELTIN			
BX	+! ALMIRALL	1.2%;0.025%	N050803 001	Jul 16, 2010

CLOBAZAM

FILM; ORAL

SYMPAZAN

	+ AQUESTIVE THERAP	5MG	N210833 001	Nov 01, 2018
	+	10MG	N210833 002	Nov 01, 2018
	+!	20MG	N210833 003	Nov 01, 2018

SUSPENSION; ORAL

CLOBAZAM

AB	AMNEAL PHARMS LLC	2.5MG/ML	A210039 001	Oct 22, 2018
AB	BIONPHARMA INC	2.5MG/ML	A208819 001	Oct 22, 2018
AB	HETERO LABS LTD III	2.5MG/ML	A209796 001	Feb 24, 2020
AB	HIKMA	2.5MG/ML	A209715 001	Oct 22, 2018
AB	LUPIN LTD	2.5MG/ML	A210546 001	Dec 28, 2018
AB	MYLAN	2.5MG/ML	A211259 001	Oct 22, 2018
AB	TEVA PHARMS USA	2.5MG/ML	A211032 001	Jan 31, 2020
AB	UPSHER SMITH LABS	2.5MG/ML	A210569 001	Oct 22, 2018
AB	VISTAPHARM	2.5MG/ML	A210746 001	Jul 10, 2019
	<u>ONFI</u>			
AB	+! LUNDBECK PHARMS LLC	2.5MG/ML	N203993 001	Dec 14, 2012

TABLET; ORAL

CLOBAZAM

AB	ALKEM LABS LTD	10MG	A212714 001	Sep 06, 2019
AB		20MG	A212714 002	Sep 06, 2019
AB	AMNEAL PHARMS CO	10MG	A209718 001	Oct 22, 2018
AB		20MG	A209718 002	Oct 22, 2018
AB	BIONPHARMA INC	10MG	A208825 001	Oct 22, 2018
AB		20MG	A208825 002	Oct 22, 2018
AB	BRECKENRIDGE	10MG	A209308 001	Oct 22, 2018
AB		20MG	A209308 002	Oct 22, 2018
AB	HETERO LABS LTD III	10MG	A209795 001	Oct 22, 2018
AB		20MG	A209795 002	Oct 22, 2018
AB	HIKMA	10MG	A208785 001	Oct 22, 2018
AB		20MG	A208785 002	Oct 22, 2018
AB	LANNETT CO INC	10MG	A212092 001	Oct 30, 2019
AB		20MG	A212092 002	Oct 30, 2019
AB	LUPIN LTD	10MG	A210545 001	Dec 14, 2018
AB		20MG	A210545 002	Dec 14, 2018
AB	MICRO LABS	10MG	A211711 001	Jan 30, 2019
AB		20MG	A211711 002	Jan 30, 2019
AB	PIRAMAL HLTHCARE UK	10MG	A209808 001	Oct 22, 2018
AB		20MG	A209808 002	Oct 22, 2018
AB	UPSHER SMITH LABS	10MG	A209687 001	Oct 22, 2018
AB		20MG	A209687 002	Oct 22, 2018
AB	ZYDUS PHARMS	10MG	A211449 001	Oct 22, 2018
AB		20MG	A211449 002	Oct 22, 2018
	<u>ONFI</u>			
AB	+ LUNDBECK PHARMS LLC	10MG	N202067 002	Oct 21, 2011
AB	+!	20MG	N202067 003	Oct 21, 2011

CLOBETASOL PROPIONATE

AEROSOL, FOAM; TOPICAL

CLOBETASOL PROPIONATE

AB1	GLENMARK PHARMS LTD	0.05%	A210809 001	Feb 15, 2019
AB1	INGENUS PHARMS LLC	0.05%	A206805 001	Jul 31, 2017
AB1	PERRIGO ISRAEL	0.05%	A077763 001	Mar 10, 2008
AB1	TARO	0.05%	A208779 001	Oct 04, 2018

OLUX

AB1	+! MYLAN	0.05%	N021142 001	May 26, 2000
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CLOBETASOL PROPIONATE

AB2	GLENMARK PHARMS LTD	0.05%	A211450 001	Sep 09, 2019
AB2	PERRIGO UK FINCO	0.05%	A201402 001	Aug 14, 2012

OLUX E

AB2	+! MYLAN	0.05%	N022013 001	Jan 12, 2007
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PRESCRIPTION DRUG PRODUCT LIST

CLOBETASOL PROPIONATE

CREAM; TOPICAL

CLOBETASOL PROPIONATE

<u>AB1</u>	ACP NIMBLE	<u>0.05%</u>	<u>A074139</u>	<u>001</u>	Aug 03, 1994
<u>AB1</u>	ALEOR DERMACEUTICALS	<u>0.05%</u>	<u>A213291</u>	<u>001</u>	Jan 27, 2020
<u>AB1</u>	AMNEAL PHARMS LLC	<u>0.05%</u>	<u>A211256</u>	<u>001</u>	Dec 26, 2018
<u>AB1</u>	FOUGERA PHARMS INC	<u>0.05%</u>	<u>A074392</u>	<u>001</u>	Sep 30, 1996
<u>AB1</u>	GLENMARK PHARMS	<u>0.05%</u>	<u>A209095</u>	<u>001</u>	May 10, 2018
<u>AB1</u>	LUPIN LTD	<u>0.05%</u>	<u>A210208</u>	<u>001</u>	Jan 30, 2018
<u>AB1</u>	MYLAN	<u>0.05%</u>	<u>A075338</u>	<u>001</u>	Feb 09, 2001
<u>AB1</u>	RISING	<u>0.05%</u>	<u>A211401</u>	<u>001</u>	Jan 11, 2019
<u>AB1</u>	TARO	<u>0.05%</u>	<u>A074249</u>	<u>001</u>	Jul 08, 1996
<u>AB1</u>	TELIGENT PHARMA INC	<u>0.05%</u>	<u>A209974</u>	<u>001</u>	Apr 17, 2018
<u>AB1</u>	TORRENT	<u>0.05%</u>	<u>A211836</u>	<u>001</u>	Dec 30, 2019
<u>AB1</u>	XIROMED	<u>0.05%</u>	<u>A210034</u>	<u>001</u>	Jun 15, 2018
<u>AB1</u>	ZYDUS PHARMS	<u>0.05%</u>	<u>A211074</u>	<u>001</u>	Oct 15, 2018

CORMAX

<u>AB1</u>	! HI TECH PHARMA	<u>0.05%</u>	<u>A074220</u>	<u>001</u>	May 16, 1997
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CLOBETASOL PROPIONATE (EMOLLIENT)

<u>AB2</u>	! FOUGERA PHARMS	<u>0.05%</u>	<u>A075430</u>	<u>001</u>	May 26, 1999
<u>AB2</u>	NOVAST LABS	<u>0.05%</u>	<u>A075733</u>	<u>001</u>	Aug 22, 2001
<u>AB2</u>	TARO	<u>0.05%</u>	<u>A075633</u>	<u>001</u>	May 17, 2000
<u>AB2</u>	TELIGENT PHARMA INC	<u>0.05%</u>	<u>A209411</u>	<u>001</u>	Aug 21, 2017

EMBELINE E

<u>AB2</u>	HI TECH PHARMA	<u>0.05%</u>	<u>A075325</u>	<u>001</u>	Dec 24, 1998
	IMPOYZ				
	+! ENCORE DERMAT	0.025%	N209483	001	Nov 28, 2017

GEL; TOPICAL

CLOBETASOL PROPIONATE

<u>AB</u>	! FOUGERA PHARMS	<u>0.05%</u>	<u>A075368</u>	<u>001</u>	Feb 15, 2000
<u>AB</u>	PERRIGO CO	<u>0.05%</u>	<u>A075027</u>	<u>001</u>	Oct 31, 1997
<u>AB</u>	TARO	<u>0.05%</u>	<u>A075279</u>	<u>001</u>	May 28, 1999
<u>AB</u>	TELIGENT PHARMA INC	<u>0.05%</u>	<u>A208881</u>	<u>001</u>	Mar 06, 2017

EMBELINE

<u>AB</u>	HI TECH PHARMA	<u>0.05%</u>	<u>A076141</u>	<u>001</u>	Apr 12, 2002
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LOTION; TOPICAL

CLOBETASOL PROPIONATE

<u>AB</u>	ACTAVIS MID ATLANTIC	<u>0.05%</u>	<u>A078223</u>	<u>001</u>	Dec 04, 2008
<u>AB</u>	CADILA	<u>0.05%</u>	<u>A205249</u>	<u>001</u>	Sep 24, 2019
<u>AB</u>	LUPIN LTD	<u>0.05%</u>	<u>A209147</u>	<u>001</u>	Sep 22, 2017
<u>AB</u>	TARO	<u>0.05%</u>	<u>A200302</u>	<u>001</u>	Jul 02, 2012
<u>AB</u>	TELIGENT PHARMA INC	<u>0.05%</u>	<u>A208667</u>	<u>001</u>	Nov 29, 2016

CLOBEX

<u>AB</u>	+! GALDERMA LABS LP	<u>0.05%</u>	<u>N021535</u>	<u>001</u>	Jul 24, 2003
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OINTMENT; TOPICAL

CLOBETASOL PROPIONATE

<u>AB</u>	ACP NIMBLE	<u>0.05%</u>	<u>A074089</u>	<u>001</u>	Feb 16, 1994
<u>AB</u>	ALEOR DERMACEUTICALS	<u>0.05%</u>	<u>A211800</u>	<u>001</u>	Mar 04, 2019
<u>AB</u>	ENCUBE	<u>0.05%</u>	<u>A211295</u>	<u>001</u>	Nov 15, 2019
<u>AB</u>	! FOUGERA PHARMS	<u>0.05%</u>	<u>A074407</u>	<u>001</u>	Feb 23, 1996
<u>AB</u>	GLENMARK PHARMS	<u>0.05%</u>	<u>A208933</u>	<u>001</u>	Mar 20, 2017
<u>AB</u>	MYLAN	<u>0.05%</u>	<u>A075057</u>	<u>001</u>	Aug 12, 1998
<u>AB</u>	NOVEL LABS INC	<u>0.05%</u>	<u>A208841</u>	<u>001</u>	May 04, 2018
<u>AB</u>	TARO	<u>0.05%</u>	<u>A074248</u>	<u>001</u>	Jul 12, 1996
<u>AB</u>	TELIGENT PHARMA INC	<u>0.05%</u>	<u>A208589</u>	<u>001</u>	Jan 23, 2019
<u>AB</u>	TORRENT	<u>0.05%</u>	<u>A212926</u>	<u>001</u>	Oct 25, 2019
<u>AB</u>	XIROMED	<u>0.05%</u>	<u>A209701</u>	<u>001</u>	Apr 17, 2018
<u>AB</u>	ZYDUS PHARMS	<u>0.05%</u>	<u>A210199</u>	<u>001</u>	Oct 27, 2017

EMBELINE

<u>AB</u>	HI TECH PHARMA	<u>0.05%</u>	<u>A074221</u>	<u>001</u>	Mar 31, 1995
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SHAMPOO; TOPICAL

CLOBETASOL PROPIONATE

<u>AB</u>	ACTAVIS MID ATLANTIC	<u>0.05%</u>	<u>A078854</u>	<u>001</u>	Jun 07, 2011
<u>AB</u>	HI TECH	<u>0.05%</u>	<u>A209871</u>	<u>001</u>	Oct 27, 2017
<u>AB</u>	PERRIGO ISRAEL	<u>0.05%</u>	<u>A090974</u>	<u>001</u>	Aug 09, 2012

CLOBEX

<u>AB</u>	+! GALDERMA LABS	<u>0.05%</u>	<u>N021644</u>	<u>001</u>	Feb 05, 2004
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PRESCRIPTION DRUG PRODUCT LIST

CLOBETASOL PROPIONATE

SOLUTION;TOPICAL

CLOBETASOL PROPIONATE

AT	ACP NIMBLE	<u>0.05%</u>	<u>A074331 001</u>	Dec 15, 1995
AT	ALEOR DERMACEUTICALS	<u>0.05%</u>	<u>A212881 001</u>	Oct 21, 2019
AT	FOUGERA PHARMS	<u>0.05%</u>	<u>A075391 001</u>	Feb 08, 1999
AT	GLENMARK PHARMS LTD	<u>0.05%</u>	<u>A210190 001</u>	Apr 18, 2018
AT	MACLEODS PHARMS LTD	<u>0.05%</u>	<u>A209361 001</u>	Oct 25, 2017
AT	NOVEL LABS INC	<u>0.05%</u>	<u>A206075 001</u>	Nov 23, 2015
AT	SAPTALIS PHARMS	<u>0.05%</u>	<u>A211494 001</u>	Oct 02, 2019
AT	TARO	<u>0.05%</u>	<u>A075224 001</u>	Nov 16, 1998
AT		<u>0.05%</u>	<u>A075363 001</u>	Dec 29, 2000
AT	TOLMAR	<u>0.05%</u>	<u>A076977 001</u>	Aug 05, 2005
AT	WOCKHARDT BIO AG	<u>0.05%</u>	<u>A075205 001</u>	Nov 13, 1998

EMBELINE

AT	! HI TECH PHARMA	<u>0.05%</u>	<u>A074222 001</u>	Dec 06, 1995
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SPRAY;TOPICAL

CLOBETASOL PROPIONATE

AT	AKORN	<u>0.05%</u>	<u>A207218 001</u>	Apr 28, 2017
AT	GLENMARK PHARMS	<u>0.05%</u>	<u>A209004 001</u>	Mar 26, 2018
AT	LUPIN LTD	<u>0.05%</u>	<u>A208125 001</u>	Mar 26, 2018
AT	PADDOCK LLC	<u>0.05%</u>	<u>A090898 001</u>	Jun 16, 2011
AT	TARO	<u>0.05%</u>	<u>A208842 001</u>	Mar 26, 2018
AT	ZYDUS PHARMS	<u>0.05%</u>	<u>A206378 001</u>	Feb 16, 2017

CLOBEX

AT	+! GALDERMA LABS LP	<u>0.05%</u>	<u>N021835 001</u>	Oct 27, 2005
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CLOCORTOLONE PIVALATE

CREAM;TOPICAL

CLODERM

+!	EPI HLTH	0.1%	<u>N017765 001</u>	
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CLOFARABINE

SOLUTION;INTRAVENOUS

CLOFARABINE

AP	ABON PHARMS LLC	<u>20MG/20ML (1MG/ML)</u>	<u>A204029 001</u>	May 09, 2017
AP	ACCORD HLTHCARE	<u>20MG/20ML (1MG/ML)</u>	<u>A212034 001</u>	Feb 22, 2019
AP	AMNEAL	<u>20MG/20ML (1MG/ML)</u>	<u>A208857 001</u>	Nov 06, 2017
AP	DR REDDYS LABS LTD	<u>20MG/20ML (1MG/ML)</u>	<u>A205375 001</u>	Nov 06, 2017
AP	GLAND PHARMA LTD	<u>20MG/20ML (1MG/ML)</u>	<u>A207831 001</u>	Oct 31, 2018
AP	INGENUS PHARMS LLC	<u>20MG/20ML (1MG/ML)</u>	<u>A210270 001</u>	Sep 14, 2018
AP	MSN	<u>20MG/20ML (1MG/ML)</u>	<u>A209775 001</u>	Dec 06, 2017
AP	MYLAN LABS LTD	<u>20MG/20ML (1MG/ML)</u>	<u>A208860 001</u>	Nov 06, 2017

CLOLAR

AP	+! GENZYME	<u>20MG/20ML (1MG/ML)</u>	<u>N021673 001</u>	Dec 28, 2004
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CLOMIPHENE CITRATE

TABLET;ORAL

CLOMIPHENE CITRATE

!	PAR PHARM	50MG	<u>A075528 001</u>	Aug 30, 1999
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CLOMIPRAMINE HYDROCHLORIDE

CAPSULE;ORAL

ANAFRANIL

AB	+! SPECGX LLC	<u>25MG</u>	<u>N019906 001</u>	Dec 29, 1989
AB	+	<u>50MG</u>	<u>N019906 002</u>	Dec 29, 1989
AB	+	<u>75MG</u>	<u>N019906 003</u>	Dec 29, 1989

CLOMIPRAMINE HYDROCHLORIDE

AB	AMNEAL PHARMS CO	<u>25MG</u>	<u>A208632 001</u>	Oct 31, 2018
AB		<u>50MG</u>	<u>A208632 002</u>	Oct 31, 2018
AB		<u>75MG</u>	<u>A208632 003</u>	Oct 31, 2018
AB	JUBILANT CADISTA	<u>25MG</u>	<u>A212218 001</u>	Oct 21, 2019
AB		<u>50MG</u>	<u>A212218 002</u>	Oct 21, 2019
AB		<u>75MG</u>	<u>A212218 003</u>	Oct 21, 2019
AB	LEADING PHARMA LLC	<u>25MG</u>	<u>A211364 001</u>	Feb 07, 2020
AB		<u>50MG</u>	<u>A211364 002</u>	Feb 07, 2020
AB		<u>75MG</u>	<u>A211364 003</u>	Feb 07, 2020
AB	LUPIN LTD	<u>25MG</u>	<u>A209294 001</u>	Nov 21, 2018
AB		<u>50MG</u>	<u>A209294 002</u>	Nov 21, 2018
AB		<u>75MG</u>	<u>A209294 003</u>	Nov 21, 2018
AB	MANKIND PHARMA	<u>25MG</u>	<u>A211767 001</u>	Apr 08, 2019
AB		<u>50MG</u>	<u>A211767 002</u>	Apr 08, 2019
AB		<u>75MG</u>	<u>A211767 003</u>	Apr 08, 2019
AB	MYLAN	<u>25MG</u>	<u>A074947 001</u>	Apr 30, 1998

PRESCRIPTION DRUG PRODUCT LIST

CLOMIPRAMINE HYDROCHLORIDE

CAPSULE;ORAL

CLOMIPRAMINE HYDROCHLORIDE

<u>AB</u>		<u>50MG</u>	<u>A074947 002</u>	Apr 30, 1998
<u>AB</u>		<u>75MG</u>	<u>A074947 003</u>	Apr 30, 1998
<u>AB</u>	SANDOZ	<u>25MG</u>	<u>A074364 001</u>	Mar 29, 1996
<u>AB</u>		<u>25MG</u>	<u>A074953 001</u>	Jun 25, 1997
<u>AB</u>		<u>50MG</u>	<u>A074364 002</u>	Mar 29, 1996
<u>AB</u>		<u>50MG</u>	<u>A074953 002</u>	Jun 25, 1997
<u>AB</u>		<u>75MG</u>	<u>A074364 003</u>	Mar 29, 1996
<u>AB</u>		<u>75MG</u>	<u>A074953 003</u>	Jun 25, 1997
<u>AB</u>	TARO	<u>25MG</u>	<u>A074694 001</u>	Dec 31, 1996
<u>AB</u>		<u>50MG</u>	<u>A074694 002</u>	Dec 31, 1996
<u>AB</u>		<u>75MG</u>	<u>A074694 003</u>	Dec 31, 1996
<u>AB</u>	ZYDUS PHARMS	<u>25MG</u>	<u>A208961 001</u>	Dec 27, 2017
<u>AB</u>		<u>50MG</u>	<u>A208961 002</u>	Dec 27, 2017
<u>AB</u>		<u>75MG</u>	<u>A208961 003</u>	Dec 27, 2017

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>	MYLAN	<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>	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
<u>AB</u>	WATSON LABS	<u>0.5MG</u>	<u>A074964 001</u>	Dec 30, 1997
<u>AB</u>		<u>1MG</u>	<u>A074964 002</u>	Dec 30, 1997
<u>AB</u>		<u>2MG</u>	<u>A074964 003</u>	Dec 30, 1997

KLONOPIN

<u>AB</u>	+	ROCHE	<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 PHARMS LTD	<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>	PAR PHARM	<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

PRESCRIPTION DRUG PRODUCT LIST

CLONIDINE

FILM, EXTENDED RELEASE;TRANSDERMAL

CATAPRES-TTS-1

<u>AB</u>	+	BOEHRINGER INGELHEIM	<u>0.1MG/24HR</u>	<u>N018891 001</u>	Oct 10, 1984
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CATAPRES-TTS-2

<u>AB</u>	+	BOEHRINGER INGELHEIM	<u>0.2MG/24HR</u>	<u>N018891 002</u>	Oct 10, 1984
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CATAPRES-TTS-3

<u>AB</u>	+	BOEHRINGER INGELHEIM	<u>0.3MG/24HR</u>	<u>N018891 003</u>	Oct 10, 1984
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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>		AVEVA	<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>		MAYNE PHARMA	<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

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
<u>AP</u>	+		<u>5MG/10ML (0.5MG/ML)</u>	<u>N020615 002</u>	Apr 27, 1999

TABLET; ORAL

CATAPRES

<u>AB</u>	+	BOEHRINGER INGELHEIM	<u>0.1MG</u>	<u>N017407 001</u>	
<u>AB</u>	+		<u>0.2MG</u>	<u>N017407 002</u>	
<u>AB</u>	+		<u>0.3MG</u>	<u>N017407 003</u>	

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>		FRONTIDA BIOPHARM	<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>		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>		MYLAN	<u>0.1MG</u>	<u>A070317 002</u>	Jul 09, 1987
<u>AB</u>			<u>0.2MG</u>	<u>A070317 003</u>	Jun 09, 1987
<u>AB</u>			<u>0.3MG</u>	<u>A070317 001</u>	Jun 09, 1987
<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>		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

PRESCRIPTION DRUG PRODUCT LIST

CLONIDINE HYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL

CLONIDINE HYDROCHLORIDE

<u>AB1</u>	AMNEAL PHARMS NY	<u>0.1MG</u>	<u>A210052</u>	<u>001</u>	Nov 20, 2017
<u>AB1</u>	ANCHEN PHARMS	<u>0.1MG</u>	<u>A202984</u>	<u>001</u>	Sep 30, 2013
<u>AB1</u>	JUBILANT GENERICS	<u>0.1MG</u>	<u>A210338</u>	<u>001</u>	Jan 29, 2018
<u>AB1</u>	LUPIN LTD	<u>0.1MG</u>	<u>A209285</u>	<u>001</u>	Oct 23, 2017
<u>AB1</u>	MAYNE PHARMA INC	<u>0.1MG</u>	<u>A210680</u>	<u>001</u>	Apr 30, 2018
<u>AB1</u>	NOVAST LABS	<u>0.1MG</u>	<u>A209675</u>	<u>001</u>	Mar 05, 2019
<u>AB1</u>	UPSHER SMITH LABS	<u>0.1MG</u>	<u>A211433</u>	<u>001</u>	Oct 12, 2018
<u>AB1</u>	XIAMEN LP PHARM CO	<u>0.1MG</u>	<u>A209757</u>	<u>001</u>	Nov 20, 2017

KAPVAY

<u>AB1</u>	+! CONCORDIA PHARMS INC	<u>0.1MG</u>	<u>N022331</u>	<u>003</u>	Sep 28, 2010
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CLONIDINE HYDROCHLORIDE

<u>AB2</u>	ACTAVIS ELIZABETH	<u>0.1MG</u>	<u>A202792</u>	<u>001</u>	May 15, 2015
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CLOPIDOGREL BISULFATE

TABLET;ORAL

CLOPIDOGREL BISULFATE

<u>AB</u>	ACCORD HLTHCARE	<u>EQ 75MG BASE</u>	<u>A202925</u>	<u>001</u>	Mar 27, 2013
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A202925</u>	<u>002</u>	Mar 27, 2013
<u>AB</u>	ACME LABS	<u>EQ 75MG BASE</u>	<u>A078004</u>	<u>001</u>	May 17, 2012
<u>AB</u>	AMNEAL PHARMS	<u>EQ 75MG BASE</u>	<u>A203751</u>	<u>001</u>	Apr 11, 2014
<u>AB</u>	APOTEX INC	<u>EQ 75MG BASE</u>	<u>A076274</u>	<u>001</u>	May 17, 2012
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A076274</u>	<u>002</u>	Mar 04, 2014
<u>AB</u>	AUROBINDO PHARMA LTD	<u>EQ 75MG BASE</u>	<u>A090540</u>	<u>001</u>	May 17, 2012
<u>AB</u>	CELLTRION	<u>EQ 75MG BASE</u>	<u>A202266</u>	<u>001</u>	Aug 14, 2012
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A202266</u>	<u>002</u>	Nov 20, 2012
<u>AB</u>	CSPC OUYI	<u>EQ 75MG BASE</u>	<u>A204359</u>	<u>001</u>	Feb 02, 2017
<u>AB</u>	DR REDDYS LABS INC	<u>EQ 75MG BASE</u>	<u>A076273</u>	<u>001</u>	Jan 14, 2008
<u>AB</u>	DR REDDYS LABS LTD	<u>EQ 300MG BASE</u>	<u>A091023</u>	<u>001</u>	May 17, 2012
<u>AB</u>	MACLEODS PHARMS LTD	<u>EQ 75MG BASE</u>	<u>A202928</u>	<u>001</u>	Feb 10, 2014
<u>AB</u>	PRINSTON INC	<u>EQ 75MG BASE</u>	<u>A206376</u>	<u>001</u>	May 07, 2018
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A206376</u>	<u>002</u>	May 07, 2018
<u>AB</u>	SCIEGEN PHARMS INC	<u>EQ 75MG BASE</u>	<u>A204165</u>	<u>001</u>	Sep 15, 2014
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A204165</u>	<u>002</u>	Sep 15, 2014
<u>AB</u>	SUN PHARM	<u>EQ 75MG BASE</u>	<u>A090494</u>	<u>001</u>	May 17, 2012
<u>AB</u>	TEVA	<u>EQ 75MG BASE</u>	<u>A076999</u>	<u>001</u>	May 17, 2012
<u>AB</u>	TORRENT PHARMS LTD	<u>EQ 75MG BASE</u>	<u>A090844</u>	<u>001</u>	May 17, 2012

PLAVIX

<u>AB</u>	+ SANOFI AVENTIS US	<u>EQ 75MG BASE</u>	<u>N020839</u>	<u>001</u>	Nov 17, 1997
<u>AB</u>	+!	<u>EQ 300MG BASE</u>	<u>N020839</u>	<u>002</u>	Sep 20, 2007

CLORAZEPATE DIPOTASSIUM

TABLET;ORAL

CLORAZEPATE DIPOTASSIUM

<u>AB</u>	MYLAN	<u>3.75MG</u>	<u>A071858</u>	<u>002</u>	Jul 17, 1987
<u>AB</u>		<u>7.5MG</u>	<u>A071858</u>	<u>003</u>	Jul 17, 1987
<u>AB</u>	!	<u>15MG</u>	<u>A071858</u>	<u>001</u>	Jul 17, 1987
<u>AB</u>	TARO	<u>3.75MG</u>	<u>A075731</u>	<u>003</u>	Apr 27, 2000
<u>AB</u>		<u>7.5MG</u>	<u>A075731</u>	<u>002</u>	Apr 27, 2000
<u>AB</u>		<u>15MG</u>	<u>A075731</u>	<u>001</u>	Apr 27, 2000

GEN-XENE

<u>AB</u>	ALRA	<u>3.75MG</u>	<u>A071787</u>	<u>001</u>	Apr 26, 1988
<u>AB</u>		<u>7.5MG</u>	<u>A071788</u>	<u>001</u>	Apr 26, 1988
<u>AB</u>		<u>15MG</u>	<u>A071789</u>	<u>001</u>	Apr 26, 1988

FRANKENE

<u>AB</u>	+ RECORDATI RARE	<u>7.5MG</u>	<u>N017105</u>	<u>007</u>	
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CLOTRIMAZOLE

CREAM;TOPICAL

CLOTRIMAZOLE

<u>AB</u>	FOUGERA PHARMS	<u>1%</u>	<u>A078338</u>	<u>001</u>	Sep 02, 2008
<u>AB</u>	GLENMARK PHARMS	<u>1%</u>	<u>A090219</u>	<u>001</u>	Aug 03, 2010
<u>AB</u>	! TARO	<u>1%</u>	<u>A072640</u>	<u>001</u>	Aug 31, 1993

SOLUTION;TOPICAL

CLOTRIMAZOLE

<u>AT</u>	NOVITIUM PHARMA	<u>1%</u>	<u>A209815</u>	<u>001</u>	Feb 14, 2019
<u>AT</u>	! TARO	<u>1%</u>	<u>A074580</u>	<u>001</u>	Jul 29, 1996
<u>AT</u>	TASMAN PHARMA	<u>1%</u>	<u>A212281</u>	<u>001</u>	Jul 25, 2019
<u>AT</u>	TEVA	<u>1%</u>	<u>A073306</u>	<u>001</u>	Feb 28, 1995

PRESCRIPTION DRUG PRODUCT LIST

CLOTRIMAZOLE

TROCHE/LOZENGE;ORAL

CLOTRIMAZOLE

<u>AB</u>	!	HIKMA	<u>10MG</u>	<u>A076387</u>	<u>001</u>	Jul 29, 2004
<u>AB</u>		PADDOCK LLC	<u>10MG</u>	<u>A076763</u>	<u>001</u>	Oct 28, 2005

CLOZAPINE

SUSPENSION;ORAL

VERSACLOZ

+	!	TASMAN PHARMA	50MG/ML	N203479	001	Feb 06, 2013
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TABLET;ORAL

CLOZAPINE

<u>AB</u>		ACCORD HLTHCARE	<u>25MG</u>	<u>A202873</u>	<u>001</u>	Nov 25, 2015
<u>AB</u>			<u>100MG</u>	<u>A202873</u>	<u>002</u>	Nov 25, 2015
<u>AB</u>		AUROBINDO PHARMA LTD	<u>25MG</u>	<u>A206433</u>	<u>001</u>	Nov 29, 2016
<u>AB</u>			<u>50MG</u>	<u>A206433</u>	<u>002</u>	Nov 29, 2016
<u>AB</u>			<u>100MG</u>	<u>A206433</u>	<u>003</u>	Nov 29, 2016
<u>AB</u>			<u>200MG</u>	<u>A206433</u>	<u>004</u>	Nov 29, 2016
<u>AB</u>		IVAX SUB TEVA PHARMS	<u>25MG</u>	<u>A074949</u>	<u>001</u>	Nov 26, 1997
<u>AB</u>			<u>50MG</u>	<u>A074949</u>	<u>004</u>	Apr 25, 2005
<u>AB</u>			<u>50MG</u>	<u>A076809</u>	<u>003</u>	Dec 16, 2005
<u>AB</u>			<u>100MG</u>	<u>A074949</u>	<u>002</u>	Nov 26, 1997
<u>AB</u>			<u>100MG</u>	<u>A076809</u>	<u>002</u>	Dec 16, 2005
<u>AB</u>			<u>200MG</u>	<u>A076809</u>	<u>001</u>	Dec 16, 2005
<u>AB</u>		MAYNE PHARMA	<u>25MG</u>	<u>A203807</u>	<u>001</u>	Sep 17, 2015
<u>AB</u>			<u>50MG</u>	<u>A203807</u>	<u>003</u>	Aug 22, 2017
<u>AB</u>			<u>100MG</u>	<u>A203807</u>	<u>002</u>	Sep 17, 2015
<u>AB</u>			<u>200MG</u>	<u>A203807</u>	<u>004</u>	Aug 22, 2017
<u>AB</u>		MYLAN	<u>25MG</u>	<u>A075417</u>	<u>001</u>	May 27, 1999
<u>AB</u>			<u>50MG</u>	<u>A075417</u>	<u>004</u>	Apr 15, 2010
<u>AB</u>			<u>100MG</u>	<u>A075417</u>	<u>002</u>	May 27, 1999
<u>AB</u>			<u>200MG</u>	<u>A075417</u>	<u>005</u>	Apr 15, 2010
<u>AB</u>		SUN PHARM INDS INC	<u>25MG</u>	<u>A075713</u>	<u>001</u>	Nov 15, 2002
<u>AB</u>			<u>50MG</u>	<u>A075713</u>	<u>003</u>	Aug 19, 2005
<u>AB</u>			<u>100MG</u>	<u>A075713</u>	<u>002</u>	Nov 15, 2002
<u>AB</u>			<u>200MG</u>	<u>A075713</u>	<u>004</u>	Nov 07, 2017

CLOZARIL

<u>AB</u>	+	HERITAGE LIFE	<u>25MG</u>	<u>N019758</u>	<u>001</u>	Sep 26, 1989
<u>AB</u>	+		<u>50MG</u>	<u>N019758</u>	<u>003</u>	May 20, 2019
<u>AB</u>	+	!	<u>100MG</u>	<u>N019758</u>	<u>002</u>	Sep 26, 1989
<u>AB</u>	+		<u>200MG</u>	<u>N019758</u>	<u>004</u>	May 20, 2019

CLOZAPINE

IVAX SUB TEVA PHARMS

A074949 003 Jul 31, 2003

TABLET, ORALLY DISINTEGRATING;ORAL

CLOZAPINE

<u>AB</u>		BARR LABS INC	<u>25MG</u>	<u>A090308</u>	<u>001</u>	Nov 25, 2015
<u>AB</u>	!		<u>100MG</u>	<u>A090308</u>	<u>002</u>	Nov 25, 2015
<u>AB</u>		MYLAN	<u>25MG</u>	<u>A201824</u>	<u>002</u>	Sep 15, 2015
<u>AB</u>			<u>100MG</u>	<u>A201824</u>	<u>003</u>	Sep 15, 2015
		BARR LABS INC	12.5MG	A090308	003	Apr 09, 2018
		TEVA PHARMS USA	150MG	A203039	001	Nov 25, 2015
			200MG	A203039	002	Nov 25, 2015

COBICISTAT

TABLET;ORAL

TYBOST

+	!	GILEAD SCIENCES INC	150MG	N203094	001	Sep 24, 2014
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COBICISTAT; DARUNAVIR

TABLET;ORAL

PREZCOBIX

+	!	JANSSEN PRODS	150MG;800MG	N205395	001	Jan 29, 2015
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COBICISTAT; DARUNAVIR; EMTRICITABINE; TENOFOVIR ALAFENAMIDE FUMARATE

TABLET;ORAL

SYMITUZA

+	!	JANSSEN PRODS	150MG;800MG;200MG;EQ 10MG BASE	N210455	001	Jul 17, 2018
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PRESCRIPTION DRUG PRODUCT LIST

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

+! GENUS LIFESCIENCES 4% N209963 001 Dec 14, 2017

NUMBRINO

CODY LABS INC 4% N209575 001 Jan 10, 2020

CODEINE PHOSPHATE; PHENYLEPHRINE HYDROCHLORIDE; PROMETHAZINE HYDROCHLORIDE

SYRUP;ORAL

PROMETH HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE W/CODEINE PHOSPHATE

AA VINTAGE 10MG/5ML;5MG/5ML;6.25MG/5ML **A040660 001** Dec 07, 2006

PROMETH VC W/ CODEINE

AA ! ACTAVIS MID 10MG/5ML;5MG/5ML;6.25MG/5ML **A088764 001** Oct 31, 1984

ATLANTIC

PROMETHAZINE HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE W/CODEINE PHOSPHATE

AA HI-TECH PHARMA CO 10MG/5ML;5MG/5ML;6.25MG/5ML **A040674 001** Dec 23, 2014

PROMETHAZINE HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE W/CODEINE PHOSPHATE

AA AMNEAL PHARMS 10MG/5ML;5MG/5ML;6.25MG/5ML **A200963 001** Aug 26, 2015

CODEINE PHOSPHATE; PROMETHAZINE HYDROCHLORIDE

SYRUP;ORAL

PROMETHAZINE HYDROCHLORIDE AND CODEINE PHOSPHATE

AA ! ACTAVIS MID 10MG/5ML;6.25MG/5ML **A088763 001** Oct 31, 1984

ATLANTIC

AA AMNEAL PHARMS 10MG/5ML;6.25MG/5ML **A200894 001** Apr 24, 2013

AA HI TECH PHARMA 10MG/5ML;6.25MG/5ML **A040151 001** Aug 26, 1997

AA NOSTRUM LABS INC 10MG/5ML;6.25MG/5ML **A090180 001** Mar 17, 2010

AA TRIS PHARMA INC 10MG/5ML;6.25MG/5ML **A200386 001** Jun 29, 2012

AA WOCKHARDT BIO AG 10MG/5ML;6.25MG/5ML **A088875 001** Dec 17, 1984

PROMETHAZINE WITH CODEINE

AA VINTAGE 10MG/5ML;6.25MG/5ML **A040650 001** Jan 31, 2006

CODEINE PHOSPHATE; PSEUDOEPHEDRINE HYDROCHLORIDE; TRIPROLIDINE HYDROCHLORIDE

SYRUP;ORAL

TRIACIN-C

! STI PHARMA LLC 10MG/5ML;30MG/5ML;1.25MG/5ML A088704 001 Mar 22, 1985

CODEINE SULFATE

TABLET;ORAL

CODEINE SULFATE

AB + HIKMA 15MG **N022402 001** Jul 16, 2009

AB + 30MG **N022402 002** Jul 16, 2009

AB +! 60MG **N022402 003** Jul 16, 2009

AB LANNETT CO INC 15MG **A203046 001** Jun 13, 2014

AB 30MG **A203046 002** Jun 13, 2014

AB 60MG **A203046 003** Jun 13, 2014

COLCHICINE

CAPSULE;ORAL

MITIGARE

+! HIKMA INTL PHARMS 0.6MG N204820 001 Sep 26, 2014

SOLUTION;ORAL

GLOPERBA

+! AVION PHARMS 0.6MG/5ML N210942 001 Jan 30, 2019

TABLET;ORAL

COLCHICINE

AB ALKEM LABS LTD 0.6MG **A211250 001** Feb 08, 2019

AB AMNEAL PHARMS 0.6MG **A204711 001** Sep 28, 2016

AB DR REDDYS 0.6MG **A209876 001** Sep 06, 2019

AB GRANULES PHARMS 0.6MG **A210425 001** Feb 05, 2020

AB MYLAN 0.6MG **A209470 001** Sep 16, 2019

AB WATSON LABS INC 0.6MG **A204461 001** Jul 31, 2019

PRESCRIPTION DRUG PRODUCT LIST

COLCHICINE

TABLET; ORAL

COLCRYS

AB	+ !	TAKEDA PHARMS USA	0.6MG	N022352	001	Jul 29, 2009
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COLCHICINE; PROBENECID

TABLET; ORAL

COL-PROBENECID

AB	!	WATSON LABS	0.5MG;500MG	A084279	001	
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PROBENECID AND COLCHICINE

AB		NOVAST LABS	0.5MG;500MG	A040618	001	May 13, 2008
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COLESEVELAM HYDROCHLORIDE

BAR, CHEWABLE; ORAL

WELCHOL

+ !	DAIICHI SANKYO INC	3.75GM		N210895	001	Apr 03, 2019
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FOR SUSPENSION; ORAL

COLESEVELAM HYDROCHLORIDE

AB		ALKEM LABS LTD	3.75GM/PACKET	A210316	001	May 06, 2019
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AB		GLENMARK PHARMS LTD	3.75GM/PACKET	A202190	002	Jul 16, 2018
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WELCHOL

AB	+ !	DAIICHI SANKYO	3.75GM/PACKET	N022362	002	Oct 02, 2009
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COLESEVELAM HYDROCHLORIDE

		GLENMARK PHARMS LTD	1.875GM/PACKET	A202190	001	Jul 16, 2018
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TABLET; ORAL

COLESEVELAM HYDROCHLORIDE

AB		ALKEM LABS LTD	625MG	A209038	001	Oct 05, 2018
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AB		BIONPHARMA INC	625MG	A208670	001	Sep 13, 2019
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AB		DR REDDYS	625MG	A210889	001	Oct 05, 2018
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AB		GLENMARK PHARMS LTD	625MG	A203480	001	May 18, 2018
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AB		IMPAX LABS INC	625MG	A091600	001	May 16, 2018
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AB		ZYDUS PHARMS	625MG	A207765	001	Oct 07, 2019
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WELCHOL

AB	+ !	DAIICHI SANKYO	625MG	N021176	001	May 26, 2000
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COLESTIPOL HYDROCHLORIDE

GRANULE; ORAL

COLESTID

AB	+	PHARMACIA UPJOHN	5GM/SCOOPFUL	N017563	003	Sep 22, 1995
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AB	+ !		5GM/PACKET	N017563	004	Sep 22, 1995
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COLESTIPOL HYDROCHLORIDE

AB		IMPAX LABS	5GM/SCOOPFUL	A077277	001	May 02, 2006
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AB			5GM/PACKET	A077277	002	May 02, 2006
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FLAVORED COLESTID

+	PHARMACIA UPJOHN	5GM/PACKET		N017563	001	
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+		5GM/SCOOPFUL		N017563	002	
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TABLET; ORAL

COLESTID

AB	+ !	PHARMACIA UPJOHN	1GM	N020222	001	Jul 19, 1994
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COLESTIPOL HYDROCHLORIDE

AB		IMPAX LABS	1GM	A077510	001	Oct 24, 2006
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COLISTIMETHATE SODIUM

INJECTABLE; INJECTION

COLISTIMETHATE SODIUM

AP		EMCURE PHARMS LTD	EQ 150MG BASE/VIAL	A202359	001	Sep 28, 2012
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AP		FRESENIUS KABI USA	EQ 150MG BASE/VIAL	A065364	001	Apr 17, 2008
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AP		NEXUS PHARMS	EQ 150MG BASE/VIAL	A065177	001	Mar 19, 2004
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AP		SAGENT PHARMS INC	EQ 150MG BASE/VIAL	A201365	001	Feb 19, 2014
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AP		XELLIA PHARMS APS	EQ 150MG BASE/VIAL	A205356	001	May 29, 2015
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AP		XGEN PHARMS	EQ 150MG BASE/VIAL	A064216	001	Feb 26, 1999
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COLY-MYCIN M

AP	+ !	PAR STERILE PRODUCTS	EQ 150MG BASE/VIAL	N050108	002	
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COLISTIN SULFATE; HYDROCORTISONE ACETATE; NEOMYCIN SULFATE; THONZONIUM BROMIDE

SUSPENSION/DROPS; OTIC

COLY-MYCIN S

+ !	ENDO PHARMS INC	EQ 3MG BASE/ML;10MG/ML;EQ 3.3MG BASE/ML;0.5MG/ML		N050356	001	
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PRESCRIPTION DRUG PRODUCT LIST

3-111 (of 453)

<u>CONIVAPTAN HYDROCHLORIDE</u>			
INJECTABLE; INTRAVENOUS			
VAPRISOL IN 5% DEXTROSE IN PLASTIC CONTAINER			
	+	CUMBERLAND PHARMS	20MG/100ML (0.2MG/ML) N021697 002 Oct 08, 2008
<u>COPANLISIB DIHYDROCHLORIDE</u>			
POWDER; INTRAVENOUS			
ALIQOPA			
	+	BAYER HEALTHCARE	60MG/VIAL N209936 001 Sep 14, 2017
<u>COPPER</u>			
INTRAUTERINE DEVICE; INTRAUTERINE			
PARAGARD T 380A			
	+	COOPERSURGICAL	309MG/COPPER N018680 001 Nov 15, 1984
<u>CORTICORELIN OVINE TRIFLUORATE</u>			
INJECTABLE; INJECTION			
ACTHREL			
	+	FERRING	EQ 0.1MG BASE/VIAL N020162 001 May 23, 1996
<u>CORTICOTROPIN</u>			
INJECTABLE; INJECTION			
H.P. ACTHAR GEL			
	+	MALLINCKRODT ARD	80 UNITS/ML N008372 008
<u>CORTISONE ACETATE</u>			
TABLET; ORAL			
CORTISONE ACETATE			
	!	HIKMA INTL PHARMS	25MG A080776 002
<u>COSYNTROPIN</u>			
INJECTABLE; INJECTION			
<u>CORTROSYN</u>			
AP	+	AMPHASTAR PHARMS INC	0.25MG/VIAL N016750 001
<u>COSYNTROPIN</u>			
AP		MYLAN INSTITUTIONAL	0.25MG/VIAL A090574 001 Dec 17, 2009
AP		SANDOZ	0.25MG/VIAL A202147 001 Jun 29, 2012
<u>CRISABOROLE</u>			
OINTMENT; TOPICAL			
EUCRISA			
	+	ANACOR PHARMS INC	2% N207695 001 Dec 14, 2016
<u>CRIZOTINIB</u>			
CAPSULE; ORAL			
XALKORI			
	+	PF PRISM CV	200MG N202570 001 Aug 26, 2011
	+		250MG N202570 002 Aug 26, 2011
<u>CROFELEMER</u>			
TABLET, DELAYED RELEASE; ORAL			
MYTESI			
	+	NAPO PHARMS INC	125MG N202292 001 Dec 31, 2012
<u>CROMOLYN SODIUM</u>			
CONCENTRATE; ORAL			
<u>CROMOLYN SODIUM</u>			
AA		AILEX PHARMS LLC	100MG/5ML A209264 001 Oct 16, 2017
AA		MICRO LABS LTD INDIA	100MG/5ML A202745 001 Apr 04, 2013
AA		RISING	100MG/5ML A202583 001 Oct 27, 2011
<u>GASTROCROM</u>			
AA	+	MYLAN SPECIALITY LP	100MG/5ML N020479 001 Feb 29, 1996
SOLUTION; INHALATION			
<u>CROMOLYN SODIUM</u>			
AN		AILEX PHARMS LLC	10MG/ML A209453 001 Oct 16, 2017
AN	!	TEVA PHARMS	10MG/ML A075271 001 Jan 18, 2000
AN		WOCKHARDT BIO AG	10MG/ML A075346 001 Oct 25, 1999
SOLUTION/DROPS; OPHTHALMIC			
<u>CROMOLYN SODIUM</u>			
AT	!	AKORN	4% A074706 001 Apr 29, 1998
AT		SANDOZ INC	4% A075282 001 Jun 16, 1999

PRESCRIPTION DRUG PRODUCT LIST

CROTAMITON

LOTION; TOPICAL

CROTAN**AT** MARNEL PHARMS **10%** **A087204 001**EURAX**AT** +! SUN PHARM INDS INC **10%** **N009112 003**CUPRIC CHLORIDE

INJECTABLE; INJECTION

CUPRIC CHLORIDE IN PLASTIC CONTAINER

+! HOSPIRA EQ 0.4MG COPPER/ML N018960 001 Jun 26, 1986

CYANOCOBALAMIN

INJECTABLE; INJECTION

CYANOCOBALAMIN**AP** +! AM REGENT **1MG/ML** **A080737 001****AP** MYLAN LABS LTD **1MG/ML** **A204829 001** Jun 05, 2017**AP** SOMERSET THERAPS **1MG/ML** **A206503 001** Dec 11, 2015

LLC

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**

INT

VIBISONE**AP** ! FRESENIUS KABI USA **1MG/ML** **A080557 003**

SPRAY, METERED; NASAL

NASCOBAL

+! ENDO PHARMS INC 0.5MG/SPRAY N021642 001 Jan 31, 2005

CYCLOBENZAPRINE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

AMRIX**AB** + TEVA PHARMS INTL **15MG** **N021777 001** Feb 01, 2007**AB** +! **30MG** **N021777 002** Feb 01, 2007CYCLOBENZAPRINE HYDROCHLORIDE**AB** APOTEX **15MG** **A206703 001** Jul 24, 2018**AB** **30MG** **A206703 002** Jul 24, 2018**AB** TWI PHARMS INC **15MG** **A091281 001** Jan 31, 2013**AB** **30MG** **A091281 002** Jan 31, 2013

TABLET; ORAL

CYCLOBENZAPRINE HYDROCHLORIDE**AB** ACTAVIS LABS FL INC **5MG** **A071611 002** Feb 03, 2006**AB** **7.5MG** **A071611 003** Feb 03, 2006**AB** **10MG** **A071611 001** May 03, 1989**AB** ANDA REPOSITORY **5MG** **A073541 002** Apr 06, 2006**AB** **10MG** **A073541 001** May 23, 1995**AB** AUROBINDO PHARMA **5MG** **A078643 001** Sep 26, 2008**AB** **10MG** **A078643 002** Sep 26, 2008**AB** INVAGEN PHARMS **5MG** **A090478 001** Jul 23, 2010**AB** **10MG** **A090478 002** Jul 23, 2010**AB** JUBILANT CADISTA **5MG** **A077563 001** Apr 19, 2006**AB** **7.5MG** **A077563 003** Aug 25, 2017**AB** **10MG** **A077563 002** Apr 19, 2006**AB** KVK TECH **5MG** **A078048 001** Feb 28, 2011**AB** **10MG** **A078048 002** Feb 28, 2011**AB** MYLAN PHARMS INC **5MG** **A073144 002** Feb 03, 2006**AB** **7.5MG** **A073144 003** Mar 25, 2013**AB** ! **10MG** **A073144 001** May 30, 1991**AB** ORIT LABS LLC **5MG** **A078218 002** Jun 19, 2015**AB** **10MG** **A078218 001** Apr 18, 2008**AB** OXFORD PHARMS **5MG** **A077209 002** Feb 03, 2006**AB** **10MG** **A077209 001** Oct 04, 2005**AB** PLIVA **10MG** **A074421 001** Sep 29, 1995**AB** PRINSTON INC **5MG** **A077797 001** Feb 28, 2007**AB** **10MG** **A077797 002** Feb 28, 2007**AB** RUBICON **5MG** **A208170 001** May 31, 2017**AB** **7.5MG** **A208170 002** May 31, 2017**AB** **10MG** **A208170 003** May 31, 2017**AB** SUN PHARM INDS LTD **5MG** **A078722 001** May 12, 2008**AB** **7.5MG** **A078722 002** May 12, 2008**AB** **10MG** **A078722 003** May 12, 2008

PRESCRIPTION DRUG PRODUCT LIST

CYCLOPENTOLATE HYDROCHLORIDE

SOLUTION/DROPS;OPHTHALMIC

AKPENTOLATE

AT AKORN **1%** **A040164 001** Jan 13, 1997

CYCLOGYL

AT ! ALCON LABS INC **0.5%** **A084109 001**

AT ! **1%** **A084110 001**

CYCLOPENTOLATE HYDROCHLORIDE

AT AKORN INC **0.5%** **A205937 001** Dec 09, 2015

PENTOLAIR

AT BAUSCH AND LOMB **1%** **A040075 001** Apr 29, 1994

CYCLOGYL

! ALCON LABS INC **2%** A084108 001

CYCLOPENTOLATE HYDROCHLORIDE; PHENYLEPHRINE HYDROCHLORIDE

SOLUTION/DROPS;OPHTHALMIC

CYCLOMYDRIL

! ALCON LABS INC **0.2%;1%** A084300 001

CYCLOPHOSPHAMIDE

CAPSULE;ORAL

CYCLOPHOSPHAMIDE

AB ANI PHARMS INC **25MG** **A207014 001** Mar 19, 2018

AB **50MG** **A207014 002** Mar 19, 2018

AB CIPLA **25MG** **A211608 001** Jan 18, 2019

AB **50MG** **A211608 002** Jan 18, 2019

AB + HIKMA **25MG** **N203856 001** Sep 16, 2013

AB +! **50MG** **N203856 002** Sep 16, 2013

AB STI PHARMA LLC **25MG** **A209872 001** May 07, 2018

AB **50MG** **A209872 002** May 07, 2018

INJECTABLE; INJECTION

CYCLOPHOSPHAMIDE

AP AMNEAL **500MG/VIAL** **A210046 001** May 25, 2018

AP **1GM/VIAL** **A210046 002** May 25, 2018

AP **2GM/VIAL** **A210046 003** May 25, 2018

AP ! BAXTER HLTHCARE **500MG/VIAL** **A040745 001** May 21, 2008

AP ! **1GM/VIAL** **A040745 002** May 21, 2008

AP ! **2GM/VIAL** **A040745 003** May 21, 2008

AP JIANGSU HENGRUI MED **500MG/VIAL** **A204555 001** Oct 31, 2014

AP **1GM/VIAL** **A204555 002** Oct 31, 2014

AP **2GM/VIAL** **A204555 003** Oct 31, 2014

CYCLOSERINE

CAPSULE;ORAL

SEROMYCIN

! PURDUE **250MG** A060593 001

CYCLOSPORINE

CAPSULE;ORAL

CYCLOSPORINE

AB1 APOTEX **25MG** **A210721 001** Jul 10, 2019

AB1 **50MG** **A210721 002** Jul 10, 2019

AB1 **100MG** **A210721 003** Jul 10, 2019

AB1 IVAX SUB TEVA **25MG** **A065110 003** Mar 29, 2005

PHARMS

AB1 **50MG** **A065110 001** Mar 29, 2005

AB1 **100MG** **A065110 002** Mar 29, 2005

AB1 MAYNE PHARMA **25MG** **A065044 002** Dec 20, 2000

AB1 **100MG** **A065044 001** Dec 20, 2000

AB1 SANDOZ **25MG** **A065017 002** Jan 13, 2000

AB1 **100MG** **A065017 001** Jan 13, 2000

GENGRAF

AB1 ABBVIE **25MG** **A065003 001** May 12, 2000

AB1 **100MG** **A065003 003** May 12, 2000

NEORAL

AB1 + NOVARTIS **25MG** **N050715 001** Jul 14, 1995

AB1 +! **100MG** **N050715 002** Jul 14, 1995

CYCLOSPORINE

AB2 APOTEX **25MG** **A065040 001** May 09, 2002

AB2 **100MG** **A065040 002** May 09, 2002

SANDIMMUNE

AB2 + NOVARTIS **25MG** **N050625 001** Mar 02, 1990

AB2 +! **100MG** **N050625 002** Mar 02, 1990

BX + **50MG** N050625 003 Nov 23, 1992

PRESCRIPTION DRUG PRODUCT LIST

CYCLOSPORINE

EMULSION;OPHTHALMIC

RESTASIS

+! ALLERGAN 0.05% N050790 001 Dec 23, 2002

RESTASIS MULTIDOSE

+! ALLERGAN 0.05% N050790 002 Oct 27, 2016

INJECTABLE; INJECTION

CYCLOSPORINE**AP** AM REGENT **50MG/ML** **A065151 001** Oct 07, 2003**AP** WEST-WARD PHARMS **50MG/ML** **A065004 001** Oct 29, 1999

INT

SANDIMMUNE**AP** +! NOVARTIS **50MG/ML** **N050573 001** Nov 14, 1983

SOLUTION;OPHTHALMIC

CEQUA

+! SUN PHARMA GLOBAL 0.09% N210913 001 Aug 14, 2018

SOLUTION;ORAL

CYCLOSPORINE**AB1** ABBVIE **100MG/ML** **A065025 001** Mar 03, 2000**AB1** IVAX SUB TEVA **100MG/ML** **A065078 001** Mar 25, 2005

PHARMS

AB1 MAYNE PHARMA **100MG/ML** **A065054 001** Dec 18, 2001NEORAL**AB1** +! NOVARTIS **100MG/ML** **N050716 001** Jul 14, 1995CYCLOSPORINE**AB2** WOCKHARDT BIO AG **100MG/ML** **A065133 001** Sep 17, 2004SANDIMMUNE**AB2** +! NOVARTIS **100MG/ML** **N050574 001** Nov 14, 1983CYPROHEPTADINE HYDROCHLORIDE

SYRUP;ORAL

CYPROHEPTADINE HYDROCHLORIDE**AA** ANDA REPOSITORY **2MG/5ML** **A204823 001** Dec 27, 2016**AA** ELYSIUM **2MG/5ML** **A209108 001** Oct 16, 2018**AA** LANNETT CO INC **2MG/5ML** **A203191 001** Jul 13, 2017**AA** ! LYNE **2MG/5ML** **A040668 001** Jun 28, 2006**AA** PHARM ASSOC **2MG/5ML** **A091295 001** Mar 28, 2013**AA** QUAGEN **2MG/5ML** **A212423 001** May 22, 2019

TABLET;ORAL

CYPROHEPTADINE HYDROCHLORIDE**AA** APEX PHARMS INC **4MG** **A207783 001** Dec 29, 2016**AA** BEXIMCO PHARMS USA **4MG** **A206676 001** Apr 12, 2019**AA** BOSCOGEN **4MG** **A040644 001** May 30, 2006**AA** ELYSIUM **4MG** **A207555 001** Jan 31, 2017**AA** ! HERITAGE PHARMA **4MG** **A087056 001****AA** MOUNTAIN **4MG** **A040537 001** Sep 30, 2003**AA** NOVAST LABS **4MG** **A205087 001** Sep 23, 2015**AA** PAR PHARM **4MG** **A087129 001****AA** STRIDES PHARMA **4MG** **A209172 001** Apr 11, 2018**AA** TWI PHARMS **4MG** **A206553 001** Nov 29, 2016**AA** ZYDUS PHARMS **4MG** **A208938 001** May 19, 2017CYSTEAMINE 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 PHARMA USA EQ 25MG BASE N203389 001 Apr 30, 2013

+! EQ 75MG BASE N203389 002 Apr 30, 2013

GRANULE;ORAL

PROCYSBI

+ HORIZON PHARMA USA EQ 75MG BASE/PACKET N213491 001 Feb 14, 2020

+! EQ 300MG BASE/PACKET N213491 002 Feb 14, 2020

CYSTEAMINE HYDROCHLORIDE

SOLUTION/DROPS;OPHTHALMIC

CYSTARAN

+! LEADIANT BIOSCI INC EQ 0.44% BASE N200740 001 Oct 02, 2012

PRESCRIPTION DRUG PRODUCT LIST

CYSTEINE HYDROCHLORIDE

SOLUTION; INTRAVENOUS

ELCYS

+	!	EXELA PHARMA SCS LLC	500MG/10ML (50MG/ML)	N210660 001	Apr 16, 2019
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NOURESS

		AVADEL LEGACY	500MG/10ML (50MG/ML)	N212535 001	Dec 13, 2019
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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	!	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	A206190 001	Nov 09, 2017
AP			100MG/ML	A205696 001	Jul 17, 2018
AP		MYLAN LABS LTD	20MG/ML	A200914 001	Dec 13, 2011
AP			20MG/ML	A200915 001	Dec 13, 2011
AP			100MG/ML	A201784 001	Jan 30, 2012
AP		WEST-WARD PHARMS INT	100MG/VIAL	A071471 001	Aug 02, 1989
AP	!		2GM/VIAL	A074245 002	Aug 31, 1994
	!		500MG/VIAL	A071472 001	Aug 02, 1989
	!		1GM/VIAL	A074245 001	Aug 31, 1994

CYTARABINE; DAUNORUBICIN

POWDER; INTRAVENOUS

VYXEOS

+	!	CELATOR PHARMS	100MG; 44MG	N209401 001	Aug 03, 2017
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DABIGATRAN ETEXILATE MESYLATE

CAPSULE; ORAL

DABIGATRAN ETEXILATE MESYLATE

AB		ALKEM LABS LTD	EQ 75MG BASE	A208040 001	Mar 11, 2020
AB			EQ 150MG BASE	A208040 002	Mar 11, 2020
PRADAXA					
AB	+	BOEHRINGER INGELHEIM	EQ 75MG BASE	N022512 001	Oct 19, 2010
AB	+	!	EQ 150MG BASE	N022512 002	Oct 19, 2010
	+		EQ 110MG BASE	N022512 003	Nov 20, 2015

DABRAFENIB MESYLATE

CAPSULE; ORAL

TAFINLAR

+		NOVARTIS	EQ 50MG BASE	N202806 001	May 29, 2013
+	!		EQ 75MG BASE	N202806 002	May 29, 2013

DACARBAZINE

INJECTABLE; INJECTION

DACARBAZINE

AP	!	FRESENIUS KABI USA	200MG/VIAL	A075371 002	Aug 27, 1999
AP		HOSPIRA	200MG/VIAL	A075940 001	Oct 18, 2001
AP		TEVA PHARMS USA	200MG/VIAL	A075259 002	Aug 27, 1998
AP	!		500MG/VIAL	A075259 001	Sep 22, 2000
AP		WEST-WARD PHARMS INT	200MG/VIAL	A075812 001	Jun 15, 2001
AP			500MG/VIAL	A075812 002	Oct 31, 2002
	!	FRESENIUS KABI USA	100MG/VIAL	A075371 001	Aug 27, 1999

DACOMITINIB

TABLET; ORAL

VIZIMPRO

+		PFIZER INC	15MG	N211288 001	Sep 27, 2018
+			30MG	N211288 002	Sep 27, 2018
+	!		45MG	N211288 003	Sep 27, 2018

DACTINOMYCIN

INJECTABLE; INJECTION

COSMEGEN

AP	+	!	RECORDATI RARE	0.5MG/VIAL	N050682 001
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DACTINOMYCIN

AP		HISUN PHARM HANGZHOU	0.5MG/VIAL	A207232 001	Jul 16, 2019
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PRESCRIPTION DRUG PRODUCT LIST

DACTINOMYCIN

INJECTABLE; INJECTION

DACTINOMYCIN

AP	MYLAN LABS LTD	0.5MG/VIAL	A203385 001	Nov 09, 2017
AP	XGEN PHARMS	0.5MG/VIAL	A203999 001	May 20, 2019

DALBAVANCIN HYDROCHLORIDE

POWDER; INTRAVENOUS

DALVANCE

+	ALLERGAN	EQ 500MG BASE/VIAL	N021883 001	May 23, 2014
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DALFAMPRIDINE

TABLET, EXTENDED RELEASE; ORAL

AMPYRA

AB	+	ACORDA	10MG	N022250 001	Jan 22, 2010
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DALFAMPRIDINE

AB		ACCORD HLTHCARE	10MG	A206863 001	Jul 11, 2018
AB		ACTAVIS LABS FL INC	10MG	A206836 001	Jan 23, 2017
AB		ALKEM LABS LTD	10MG	A206765 001	Jul 30, 2018
AB		AUROBINDO PHARMA LTD	10MG	A206811 001	Jan 23, 2017
AB		HIKMA	10MG	A206646 001	Oct 24, 2018
AB		MICRO LABS	10MG	A210158 001	Mar 11, 2019
AB		SUN PHARM	10MG	A208292 001	May 21, 2019

DALFOPRISTIN; QUINUPRISTIN

INJECTABLE; INTRAVENOUS

SYNERCID

+	KING PHARMS	350MG/VIAL; 150MG/VIAL	N050748 001	Sep 21, 1999
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DALTEPARIN SODIUM

INJECTABLE; SUBCUTANEOUS

FRAGMIN

+	PFIZER INC	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
+		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

AB		BARR	50MG	A074582 003	May 29, 1998
AB			100MG	A074582 002	May 29, 1998
AB	!		200MG	A074582 001	Aug 09, 1996
AB		LANNETT CO INC	50MG	A077246 002	Apr 19, 2007
AB			100MG	A077246 003	Apr 19, 2007
AB			200MG	A077246 001	Sep 28, 2005

DANTROLENE SODIUM

CAPSULE; ORAL

DANTRIMUM

AB	+	PAR STERILE PRODUCTS	25MG	N017443 001	
AB	+		50MG	N017443 003	
AB	+		100MG	N017443 002	

DANTROLENE SODIUM

AB		ELITE LABS INC	25MG	A076686 001	Oct 24, 2005
AB			50MG	A076686 002	Oct 24, 2005
AB			100MG	A076686 003	Oct 24, 2005
AB		IMPAX LABS	25MG	A076856 001	Mar 01, 2005
AB			50MG	A076856 002	Mar 01, 2005
AB			100MG	A076856 003	Mar 01, 2005

FOR SUSPENSION; INTRAVENOUS

RYANODEX

+	EAGLE PHARMS	250MG/VIAL	N205579 001	Jul 22, 2014
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INJECTABLE; INJECTION

DANTRIMUM

AP	+	PAR STERILE PRODUCTS	20MG/VIAL	N018264 001	
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DANTROLENE SODIUM

AP		HIKMA PHARMS	20MG/VIAL	A204762 001	Jun 19, 2017
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PRESCRIPTION DRUG PRODUCT LISTDANTROLENE SODIUM

INJECTABLE; INJECTION

REVONTO

AP	US WORLDMEDS	20MG/VIAL	A078378 001	Jul 24, 2007
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DAPAGLIFLOZIN

TABLET; ORAL

FARXIGA

+	ASTRAZENECA AB	5MG	N202293 001	Jan 08, 2014
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+	!	10MG	N202293 002	Jan 08, 2014
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DAPAGLIFLOZIN; METFORMIN HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL

XIGDUO XR

+	ASTRAZENECA AB	2.5MG; 1GM	N205649 005	Jul 28, 2017
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+		5MG; 500MG	N205649 001	Oct 29, 2014
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+		5MG; 1GM	N205649 002	Oct 29, 2014
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+		10MG; 500MG	N205649 003	Oct 29, 2014
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+	!	10MG; 1GM	N205649 004	Oct 29, 2014
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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
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+		5MG; 1GM; EQ 2.5MG BASE	N210874 002	May 02, 2019
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+		5MG; 1GM; EQ 5MG BASE	N210874 003	May 02, 2019
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+	!	10MG; 1GM; EQ 5MG BASE	N210874 004	May 02, 2019
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DAPAGLIFLOZIN; SAXAGLIPTIN HYDROCHLORIDE

TABLET; ORAL

QTERN

+	ASTRAZENECA AB	5MG; EQ 5MG BASE	N209091 002	May 02, 2019
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+	!	10MG; EQ 5MG BASE	N209091 001	Feb 27, 2017
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DAPSONE

GEL; TOPICAL

ACZONE

AB	+	ALLERGAN	5%	N021794 001	Jul 07, 2005
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DAPSONE

AB		TARO	5%	A209506 001	Oct 16, 2017
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AB		TARO PHARMS	7.5%	A210191 001	Jun 26, 2019
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ACZONE

+	!	ALMIRALL	7.5%	N207154 001	Feb 24, 2016
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TABLET; ORAL

DAPSONE

AB		ACTAVIS LLC	25MG	A204380 001	Mar 23, 2017
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AB			100MG	A204380 002	Mar 23, 2017
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AB		ALVOGEN	25MG	A205429 001	Jan 07, 2016
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AB			100MG	A205429 002	Jan 07, 2016
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AB	!	JACOBUS	25MG	A086841 001	
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AB	!		100MG	A086842 001	
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AB		NOSTRUM LABS INC	25MG	A203887 001	May 06, 2016
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AB			100MG	A203887 002	May 06, 2016
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AB		NOVITIUM PHARMA	25MG	A206505 001	Dec 01, 2016
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AB			100MG	A206505 002	Dec 01, 2016
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AB		SAI LIFE SCIENCES	100MG	A207165 001	May 08, 2019
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AB		VIRTUS PHARMS	25MG	A204074 001	May 10, 2016
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AB			100MG	A204074 002	May 10, 2016
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DAPTOMYCIN

POWDER; INTRAVENOUS

CUBICIN

AP	+	CUBIST PHARMS LLC	500MG/VIAL	N021572 002	Sep 12, 2003
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DAPTOMYCIN

AP		ACCORD HLTHCARE	500MG/VIAL	A211961 001	Jun 24, 2019
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AP		BE PHARMS	500MG/VIAL	A212513 001	Jun 26, 2019
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AP		DR REDDYS	500MG/VIAL	A208375 001	May 01, 2019
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AP		FRESENIUS KABI USA	500MG/VIAL	A206077 001	Apr 11, 2018
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AP		HOSPIRA INC	500MG/VIAL	A202857 001	Sep 12, 2014
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AP		JIANGSU HENGRUI MED	500MG/VIAL	A212022 001	Aug 22, 2019
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AP		MYLAN LABS LTD	500MG/VIAL	A205037 001	Jun 05, 2018
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AP		SAGENT PHARMS INC	500MG/VIAL	A207104 001	Nov 15, 2019
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AP		TEVA PHARMS USA	500MG/VIAL	A091039 001	Mar 25, 2016
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AP		XELLIA PHARMS APS	500MG/VIAL	A206005 001	Jun 15, 2016
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PRESCRIPTION DRUG PRODUCT LIST

DAPTOMYCIN

POWDER; INTRAVENOUS
CUBICIN RF

+! CUBIST PHARMS LLC 500MG/VIAL N021572 003 Jul 06, 2016
POWDER; IV (INFUSION)

DAPTOMYCIN

AP ACCORD HLTHCARE **350MG/VIAL** **A212667 001** Jul 12, 2019
AP +! SAGENT PHARMS INC **350MG/VIAL** **N208385 001** Sep 12, 2017
+! XELLIA PHARMS APS 350MG/VIAL N209949 001 Oct 20, 2017

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, 2017

DARIFENACIN HYDROBROMIDE

AB ALAN LABS INC **EQ 7.5MG BASE** **A209571 002** Oct 22, 2019
AB **EQ 15MG BASE** **A209571 001** Oct 22, 2019
AB ALEMBIC PHARMS LTD **EQ 7.5MG BASE** **A207681 001** Dec 08, 2017
AB **EQ 15MG BASE** **A207681 002** Dec 08, 2017
AB AUROBINDO PHARMA LTD **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 JUBILANT GENERICS **EQ 7.5MG BASE** **A205550 001** Oct 12, 2016
AB **EQ 15MG BASE** **A205550 002** Oct 12, 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, 2016

ENABLEX

AB + APIL **EQ 7.5MG BASE** **N021513 001** Dec 22, 2004
AB +! **EQ 15MG BASE** **N021513 002** Dec 22, 2004

DAROLUTAMIDE

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 TEVA PHARMS USA **600MG** **A202118 001** Nov 21, 2017

PREZISTA

AB + JANSSEN PRODS **600MG** **N021976 002** Feb 25, 2008
+ 75MG N021976 004 Dec 18, 2008
+ 150MG N021976 005 Dec 18, 2008
+! 800MG N021976 006 Nov 09, 2012

DASABUVIR SODIUM; OMBITASVIR; PARITAPREVIR; RITONAVIR

TABLET, TABLET; ORAL

VIEKIRA PAK (COPACKAGED)

+! ABBEVIE INC 250.0MG; 12.5MG; 75.0MG; EQ 250MG N206619 001 Dec 19, 2014
BASE, N/A, N/A, N/A; N/A, 12.5MG, 75MG, 50MG

DASATINIB

TABLET; ORAL

SPRYCEL

+ BRISTOL MYERS 20MG N021986 001 Jun 28, 2006
SQUIBB
+ 50MG N021986 002 Jun 28, 2006
+ 70MG N021986 003 Jun 28, 2006
+ 80MG N021986 005 Oct 28, 2010
+! 100MG N021986 004 May 30, 2008
+ 140MG N021986 006 Oct 28, 2010

PRESCRIPTION DRUG PRODUCT LIST

DAUNORUBICIN HYDROCHLORIDE

INJECTABLE; INJECTION

CERUBIDINE

<u>AP</u>	!	WEST-WARD PHARMS INT	<u>EQ 20MG BASE/VIAL</u>	<u>A064103</u>	<u>001</u>	Feb 03, 1995
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DAUNORUBICIN HYDROCHLORIDE

<u>AP</u>		FRESENIUS KABI USA	<u>EQ 20MG BASE/VIAL</u>	<u>A065000</u>	<u>001</u>	May 25, 1999
<u>AP</u>		HISUN PHARM HANGZHOU	<u>EQ 5MG BASE/ML</u>	<u>A208759</u>	<u>001</u>	Apr 12, 2019
<u>AP</u>		TEVA PHARMS USA	<u>EQ 5MG BASE/ML</u>	<u>A065035</u>	<u>001</u>	Jan 24, 2000
<u>AP</u>	+	WEST-WARD PHARMS INT	<u>EQ 5MG BASE/ML</u>	<u>N050731</u>	<u>001</u>	Jan 30, 1998
		FRESENIUS KABI USA	EQ 5MG BASE/VIAL	A065034	001	Nov 20, 2001

DECITABINE

INJECTABLE; INTRAVENOUS

DACOGEN

<u>AP</u>	+	OTSUKA PHARM CO LTD	<u>50MG/VIAL</u>	<u>N021790</u>	<u>001</u>	May 02, 2006
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DECITABINE

<u>AP</u>		ACCORD HLTHCARE	<u>50MG/VIAL</u>	<u>A203475</u>	<u>001</u>	Feb 27, 2017
<u>AP</u>		CHEMI SPA	<u>50MG/VIAL</u>	<u>A206033</u>	<u>001</u>	Sep 22, 2017
<u>AP</u>		CIPLA	<u>50MG/VIAL</u>	<u>A208601</u>	<u>001</u>	Nov 16, 2017
<u>AP</u>		DR REDDYS	<u>50MG/VIAL</u>	<u>A203131</u>	<u>001</u>	Jul 11, 2013
<u>AP</u>		INGENUS PHARMS LLC	<u>50MG/VIAL</u>	<u>A210984</u>	<u>001</u>	Sep 16, 2019
<u>AP</u>		LUPIN LTD	<u>50MG/VIAL</u>	<u>A210756</u>	<u>001</u>	Nov 09, 2018
<u>AP</u>		MSN	<u>50MG/VIAL</u>	<u>A212265</u>	<u>001</u>	Aug 28, 2019
<u>AP</u>		PHARMASCIENCE INC	<u>50MG/VIAL</u>	<u>A204607</u>	<u>001</u>	May 31, 2017
<u>AP</u>		SAGENT PHARMS INC	<u>50MG/VIAL</u>	<u>A207100</u>	<u>001</u>	Mar 16, 2018
<u>AP</u>		SANDOZ INC	<u>50MG/VIAL</u>	<u>A202969</u>	<u>001</u>	Aug 28, 2014
<u>AP</u>		WOCKHARDT BIO AG	<u>50MG/VIAL</u>	<u>A209056</u>	<u>001</u>	Apr 09, 2019

POWDER; INTRAVENOUS

DECITABINE

	+	SUN PHARMA GLOBAL	50MG/VIAL	N205582	001	Jan 28, 2014
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DEFERASIROX

GRANULE; ORAL

JADENU SPRINKLE

	+	NOVARTIS	90MG	N207968	001	May 18, 2017
	+		180MG	N207968	002	May 18, 2017
	+		360MG	N207968	003	May 18, 2017

TABLET; ORAL

DEFERASIROX

<u>AB</u>		ACTAVIS ELIZABETH	<u>90MG</u>	<u>A208697</u>	<u>001</u>	Dec 13, 2019
<u>AB</u>			<u>180MG</u>	<u>A208697</u>	<u>002</u>	Dec 13, 2019
<u>AB</u>			<u>360MG</u>	<u>A208697</u>	<u>003</u>	Dec 13, 2019
<u>AB</u>		ALEMBIC PHARMS LTD	<u>90MG</u>	<u>A211824</u>	<u>001</u>	Nov 20, 2019
<u>AB</u>			<u>360MG</u>	<u>A211824</u>	<u>002</u>	Nov 20, 2019
<u>AB</u>		AMNEAL	<u>90MG</u>	<u>A210727</u>	<u>001</u>	Dec 27, 2019
<u>AB</u>			<u>360MG</u>	<u>A210727</u>	<u>002</u>	Dec 27, 2019
<u>AB</u>		CIPLA	<u>90MG</u>	<u>A211852</u>	<u>001</u>	Feb 11, 2020
<u>AB</u>			<u>360MG</u>	<u>A211852</u>	<u>002</u>	Feb 11, 2020
<u>AB</u>		MSN	<u>90MG</u>	<u>A210945</u>	<u>001</u>	Nov 20, 2019
<u>AB</u>			<u>360MG</u>	<u>A210945</u>	<u>002</u>	Nov 20, 2019
<u>AB</u>		PIRAMAL HLTHCARE UK	<u>90MG</u>	<u>A212995</u>	<u>001</u>	Dec 30, 2019
<u>AB</u>			<u>360MG</u>	<u>A212995</u>	<u>002</u>	Dec 30, 2019
<u>AB</u>		SUN PHARM	<u>90MG</u>	<u>A211641</u>	<u>001</u>	Jan 02, 2020
<u>AB</u>			<u>360MG</u>	<u>A211641</u>	<u>002</u>	Jan 02, 2020
<u>AB</u>		TEVA PHARMS USA	<u>90MG</u>	<u>A209223</u>	<u>001</u>	Nov 25, 2019
<u>AB</u>			<u>360MG</u>	<u>A209223</u>	<u>002</u>	Nov 25, 2019
<u>AB</u>		ZYDUS PHARMS	<u>90MG</u>	<u>A211383</u>	<u>001</u>	Nov 20, 2019
<u>AB</u>			<u>360MG</u>	<u>A211383</u>	<u>002</u>	Nov 20, 2019
<u>AB</u>	+	NOVARTIS PHARMS CORP	<u>90MG</u>	<u>N206910</u>	<u>001</u>	Mar 30, 2015
<u>AB</u>	+		<u>180MG</u>	<u>N206910</u>	<u>002</u>	Mar 30, 2015
<u>AB</u>	+		<u>360MG</u>	<u>N206910</u>	<u>003</u>	Mar 30, 2015

TABLET, FOR SUSPENSION; ORAL

DEFERASIROX

<u>AB</u>		ACTAVIS ELIZABETH	<u>125MG</u>	<u>A203560</u>	<u>001</u>	Jan 26, 2016
<u>AB</u>			<u>250MG</u>	<u>A203560</u>	<u>002</u>	Jan 26, 2016
<u>AB</u>			<u>500MG</u>	<u>A203560</u>	<u>003</u>	Jan 26, 2016
<u>AB</u>		ALEMBIC PHARMS LTD	<u>125MG</u>	<u>A210060</u>	<u>001</u>	Nov 20, 2019
<u>AB</u>			<u>250MG</u>	<u>A210060</u>	<u>002</u>	Nov 20, 2019
<u>AB</u>			<u>500MG</u>	<u>A210060</u>	<u>003</u>	Nov 20, 2019

PRESCRIPTION DRUG PRODUCT LIST

DEFERASIROX

TABLET, FOR SUSPENSION;ORAL

DEFERASIROX

<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 INC	<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
<u>AB</u>	ICHNOS	<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>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

+	APOPHARMA INC	80MG/ML	N208030 002	Apr 20, 2018
+	!	100MG/ML	N208030 001	Sep 09, 2015

TABLET;ORAL

FERRIPROX

+	!	APOPHARMA INC	500MG	N021825 001	Oct 14, 2011
+			1GM	N021825 002	Jul 25, 2019

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>	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>AP</u>	WEST-WARD PHARMS INT	<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>DESFERAL</u>					
<u>AP</u>	+	NOVARTIS	<u>500MG/VIAL</u>	<u>N016267 001</u>	
<u>AP</u>	+		<u>2GM/VIAL</u>	<u>N016267 002</u>	May 25, 2000

DEFIBROTIDE SODIUM

SOLUTION;INTRAVENOUS

DEFITELIO

+	!	JAZZ PHARMS INC	200MG/2.5ML (80MG/ML)	N208114 001	Mar 30, 2016
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DEFLAZACORT

SUSPENSION;ORAL

EMFLAZA

+	!	PTC THERAP	22.75MG/ML	N208685 001	Feb 09, 2017
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TABLET;ORAL

EMFLAZA

+		PTC THERAP	6MG	N208684 001	Feb 09, 2017
+			18MG	N208684 002	Feb 09, 2017
+			30MG	N208684 003	Feb 09, 2017
+	!		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

PRESCRIPTION DRUG PRODUCT LIST

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

DELAVIRDINE MESYLATE

TABLET; ORAL

RESCRIPTOR

+! VIIV HLTHCARE

200MG

N020705 002 Jul 14, 1999

DEMECLOCYCLINE HYDROCHLORIDE

TABLET; ORAL

DEMECLOCYCLINE HYDROCHLORIDE

AB	AKORN	150MG	<u>A065389 001</u>	Dec 01, 2008
AB		300MG	<u>A065389 002</u>	Dec 01, 2008
AB	AMNEAL PHARM	150MG	<u>A065425 001</u>	Feb 27, 2008
AB	!	300MG	<u>A065425 002</u>	Feb 27, 2008
AB	BARR	150MG	<u>A065171 001</u>	Dec 13, 2004
AB		300MG	<u>A065171 002</u>	Dec 13, 2004
AB	EPIC PHARMA LLC	150MG	<u>A065447 001</u>	Aug 18, 2015
AB		300MG	<u>A065447 002</u>	Aug 18, 2015

DEOXYCHOLIC ACID

SOLUTION; SUBCUTANEOUS

KYBELLA

+! KYTHERA BIOPHARMS

20MG/2ML (10MG/ML)

N206333 001 Apr 29, 2015

DESFLURANE

LIQUID; INHALATION

DESFLURANE

AN	SHANGHAI HENGRUI	100%	<u>A208234 001</u>	Feb 26, 2018
	SUPRANE			
AN	+! BAXTER HLTHCARE	100%	<u>N020118 001</u>	Sep 18, 1992

DESIPRAMINE HYDROCHLORIDE

TABLET; ORAL

DESIPRAMINE HYDROCHLORIDE

AB	ACTAVIS TOTOWA	10MG	<u>A074430 001</u>	Feb 09, 1996
AB		25MG	<u>A071601 001</u>	Jun 05, 1987
AB		50MG	<u>A071588 001</u>	Jun 05, 1987
AB		75MG	<u>A071602 001</u>	Oct 05, 1987
AB		100MG	<u>A071766 001</u>	Oct 05, 1987
AB		150MG	<u>A074430 002</u>	Feb 09, 1996
AB	AMNEAL PHARMS CO	10MG	<u>A208105 001</u>	Mar 17, 2016
AB		25MG	<u>A208105 002</u>	Mar 17, 2016
AB		50MG	<u>A208105 003</u>	Mar 17, 2016
AB		75MG	<u>A208105 004</u>	Mar 17, 2016
AB		100MG	<u>A208105 005</u>	Mar 17, 2016
AB		150MG	<u>A208105 006</u>	Mar 17, 2016
AB	ANI PHARMS INC	10MG	<u>A205153 001</u>	Oct 28, 2016
AB		25MG	<u>A205153 002</u>	Oct 28, 2016
AB		50MG	<u>A205153 003</u>	Oct 28, 2016
AB		75MG	<u>A205153 004</u>	Oct 28, 2016
AB		100MG	<u>A205153 005</u>	Oct 28, 2016
AB		150MG	<u>A205153 006</u>	Oct 28, 2016
AB	HERITAGE PHARMS INC	10MG	<u>A207433 001</u>	May 05, 2016
AB		25MG	<u>A207433 002</u>	May 05, 2016
AB		50MG	<u>A207433 003</u>	May 05, 2016
AB		75MG	<u>A207433 004</u>	May 05, 2016
AB		100MG	<u>A207433 005</u>	May 05, 2016
AB		150MG	<u>A207433 006</u>	May 05, 2016
AB	NOVAST LABS	10MG	<u>A204963 001</u>	Dec 26, 2017
AB		25MG	<u>A204963 002</u>	Dec 26, 2017
AB		50MG	<u>A204963 003</u>	Dec 26, 2017
AB		75MG	<u>A204963 004</u>	Dec 26, 2017
AB		100MG	<u>A204963 005</u>	Dec 26, 2017
AB		150MG	<u>A204963 006</u>	Dec 26, 2017
AB	SANDOZ	10MG	<u>A072099 001</u>	May 24, 1988
AB		25MG	<u>A072100 001</u>	May 24, 1988
AB		50MG	<u>A072101 001</u>	May 24, 1988
AB		75MG	<u>A072102 001</u>	Jun 20, 1988
AB		100MG	<u>A072103 001</u>	Jun 20, 1988

PRESCRIPTION DRUG PRODUCT LIST

DESIPRAMINE HYDROCHLORIDE

TABLET;ORAL

DESIPRAMINE HYDROCHLORIDE

<u>AB</u>		<u>150MG</u>	<u>A072104 001</u>	Jun 20, 1988
	<u>NORPRAMIN</u>			
<u>AB</u>	+	US PHARM HOLDINGS	<u>10MG</u>	<u>N014399 007</u> Feb 11, 1982
<u>AB</u>	+		<u>25MG</u>	<u>N014399 001</u>
<u>AB</u>	+		<u>50MG</u>	<u>N014399 003</u>
<u>AB</u>	+		<u>75MG</u>	<u>N014399 004</u>
<u>AB</u>	+		<u>100MG</u>	<u>N014399 005</u>
<u>AB</u>	+		<u>150MG</u>	<u>N014399 006</u>

DESLORATADINE

SOLUTION;ORAL

CLARINEX

<u>AA</u>	+	MERCK SHARP DOHME	<u>0.5MG/ML</u>	<u>N021300 001</u> Sep 01, 2004
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DESLORATADINE

<u>AA</u>		TARO	<u>0.5MG/ML</u>	<u>A202936 001</u> May 26, 2016
<u>AA</u>		TARO PHARM INDS LTD	<u>0.5MG/ML</u>	<u>A202592 001</u> Jun 30, 2015

TABLET;ORAL

CLARINEX

<u>AB</u>	+	MERCK SHARP DOHME	<u>5MG</u>	<u>N021165 001</u> Dec 21, 2001
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DESLORATADINE

<u>AB</u>		BELCHER PHARMS	<u>5MG</u>	<u>A078355 001</u> Apr 19, 2012
<u>AB</u>		DR REDDYS LABS LTD	<u>5MG</u>	<u>A078365 001</u> Mar 08, 2011
<u>AB</u>		LUPIN PHARMS	<u>5MG</u>	<u>A078352 001</u> Oct 25, 2010
<u>AB</u>		ORCHID HLTHCARE	<u>5MG</u>	<u>A078357 001</u> Feb 19, 2010
<u>AB</u>		PERRIGO	<u>5MG</u>	<u>A078361 001</u> Dec 22, 2011
<u>AB</u>		SANDOZ	<u>5MG</u>	<u>A078364 001</u> Dec 03, 2010
<u>AB</u>		SUN PHARM INDS	<u>5MG</u>	<u>A078359 001</u> Nov 16, 2010

TABLET, ORALLY DISINTEGRATING;ORAL

CLARINEX

<u>AB</u>	+	MERCK SHARP DOHME	<u>2.5MG</u>	<u>N021312 002</u> Jul 14, 2005
<u>AB</u>	+		<u>5MG</u>	<u>N021312 001</u> Jun 26, 2002

DESLORATADINE

<u>AB</u>		REDDYS	<u>2.5MG</u>	<u>A078367 001</u> Jul 12, 2010
<u>AB</u>			<u>5MG</u>	<u>A078367 002</u> Jul 12, 2010

DESLORATADINE; PSEUDOEPHEDRINE SULFATE

TABLET, EXTENDED RELEASE;ORAL

CLARINEX D 24 HOUR

<u>AB</u>	+	MERCK SHARP DOHME	<u>5MG;240MG</u>	<u>N021605 001</u> Mar 03, 2005
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DESLORATADINE AND PSEUDOEPHEDRINE SULFATE 24 HOUR

<u>AB</u>		DR REDDYS LABS LTD	<u>5MG;240MG</u>	<u>A078366 001</u> Apr 26, 2011
		CLARINEX-D 12 HOUR		
	+	MERCK SHARP DOHME	<u>2.5MG;120MG</u>	<u>N021313 001</u> Feb 01, 2006

DESMOPRESSIN ACETATE

INJECTABLE;INJECTION

DDAVP

<u>AP</u>	+	FERRING PHARMS INC	<u>0.004MG/ML</u>	<u>N018938 001</u> Mar 30, 1984
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DESMOPRESSIN ACETATE

<u>AP</u>		AM REGENT	<u>0.004MG/ML</u>	<u>A091374 001</u> Feb 14, 2019
<u>AP</u>		SAGENT PHARMS INC	<u>0.004MG/ML</u>	<u>A204695 001</u> Aug 22, 2017
<u>AP</u>			<u>0.004MG/ML</u>	<u>A204751 001</u> Aug 22, 2017
<u>AP</u>		SUN PHARM INDS LTD	<u>0.004MG/ML</u>	<u>A091280 001</u> Jan 25, 2013
<u>AP</u>		TEVA PHARMS USA	<u>0.004MG/ML</u>	<u>A074888 001</u> Oct 15, 1997
<u>AP</u>		UBI	<u>0.004MG/ML</u>	<u>A210218 001</u> Feb 14, 2020

SOLUTION;NASAL

DDAVP

	+	FERRING PHARMS INC	<u>0.01%</u>	<u>N017922 001</u>
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SPRAY, METERED;NASAL

DDAVP (NEEDS NO REFRIGERATION)

<u>AB</u>	+	FERRING PHARMS INC	<u>0.01MG/SPRAY</u>	<u>N017922 003</u> Aug 07, 1996
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DESMOPRESSIN ACETATE

<u>AB</u>	!	BAUSCH AND LOMB	<u>0.01MG/SPRAY</u>	<u>A074830 001</u> Jan 25, 1999
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DESMOPRESSIN ACETATE (NEEDS NO REFRIGERATION)

<u>AB</u>		APOTEX INC	<u>0.01MG/SPRAY</u>	<u>A076703 001</u> Jan 27, 2005
<u>AB</u>		SUN PHARM	<u>0.01MG/SPRAY</u>	<u>A078271 001</u> Dec 23, 2013
<u>AB</u>		ZYDUS PHARMS	<u>0.01MG/SPRAY</u>	<u>A091345 001</u> Oct 03, 2017

MINIRIN

<u>AB</u>	+	FERRING	<u>0.01MG/SPRAY</u>	<u>N021333 001</u> Sep 16, 2002
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PRESCRIPTION DRUG PRODUCT LIST

DESMOPRESSIN ACETATE

SPRAY, METERED;NASAL

STIMATE (NEEDS NO REFRIGERATION)

+! FERRING PHARMS INC 0.15MG/SPRAY

N020355 002 Oct 24, 2007

TABLET;ORAL

DDAVP

AB	+	FERRING PHARMS INC	<u>0.1MG</u>	<u>N019955</u>	<u>001</u>	Sep 06, 1995
AB	+	!	<u>0.2MG</u>	<u>N019955</u>	<u>002</u>	Sep 06, 1995

DESMOPRESSIN ACETATE

AB		ABHAI LLC	<u>0.1MG</u>	<u>A210371</u>	<u>001</u>	Jan 28, 2019
AB			<u>0.2MG</u>	<u>A210371</u>	<u>002</u>	Jan 28, 2019
AB		ACTAVIS LABS FL INC	<u>0.1MG</u>	<u>A076470</u>	<u>001</u>	Jul 01, 2005
AB			<u>0.2MG</u>	<u>A076470</u>	<u>002</u>	Jul 01, 2005
AB		APOTEX INC	<u>0.1MG</u>	<u>A077414</u>	<u>001</u>	Mar 07, 2006
AB			<u>0.2MG</u>	<u>A077414</u>	<u>002</u>	Mar 07, 2006
AB		GLENMARK PHARMS LTD	<u>0.1MG</u>	<u>A201831</u>	<u>001</u>	May 28, 2015
AB			<u>0.2MG</u>	<u>A201831</u>	<u>002</u>	May 28, 2015
AB		HERITAGE PHARMA	<u>0.1MG</u>	<u>A207880</u>	<u>001</u>	May 26, 2017
AB			<u>0.2MG</u>	<u>A207880</u>	<u>002</u>	May 26, 2017
AB		IMPAX LABS INC	<u>0.1MG</u>	<u>A077122</u>	<u>001</u>	Jan 25, 2006
AB			<u>0.2MG</u>	<u>A077122</u>	<u>002</u>	Jan 25, 2006
AB		MYLAN	<u>0.1MG</u>	<u>A200653</u>	<u>001</u>	Jun 27, 2014
AB			<u>0.2MG</u>	<u>A200653</u>	<u>002</u>	Jun 27, 2014
AB		NOVAST LABS	<u>0.1MG</u>	<u>A208357</u>	<u>001</u>	Jun 06, 2019
AB			<u>0.2MG</u>	<u>A208357</u>	<u>002</u>	Jun 06, 2019

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-28

BEKYREE

AB		LUPIN LTD	<u>0.15MG,N/A;0.02MG,0.01MG</u>	<u>A202226</u>	<u>001</u>	Aug 12, 2015
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CYCLESSA

AB	+	!	<u>0.1MG,0.125MG,0.15MG;0.025MG,0.025MG,0.025MG</u>	<u>N021090</u>	<u>001</u>	Dec 20, 2000
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DESOGESTREL AND ETHINYL ESTRADIOL

AB		ACCORD HLTHCARE	<u>0.15MG,N/A;0.02MG,0.01MG</u>	<u>A209170</u>	<u>001</u>	Jun 05, 2017
AB		AUROBINDO PHARMA LTD	<u>0.15MG,N/A;0.02MG,0.01MG</u>	<u>A206853</u>	<u>001</u>	Mar 22, 2017
AB	!	DURAMED PHARMS BARR	<u>0.15MG;0.03MG</u>	<u>A075256</u>	<u>002</u>	Aug 12, 1999
AB		MAYNE PHARMA	<u>0.15MG,N/A;0.02MG,0.01MG</u>	<u>A076916</u>	<u>001</u>	Dec 29, 2008
AB			<u>0.1MG,0.125MG,0.15MG;0.025MG,0.025MG,0.025MG</u>	<u>A077182</u>	<u>001</u>	Jan 24, 2006
AB		MYLAN LABS LTD	<u>0.15MG,N/A;0.02MG,0.01MG</u>	<u>A202296</u>	<u>001</u>	Aug 30, 2013
AB			<u>0.15MG;0.03MG</u>	<u>A202085</u>	<u>001</u>	May 20, 2015
AB		NOVAST LABS	<u>0.15MG;0.03MG</u>	<u>A091234</u>	<u>001</u>	Jul 12, 2013
AB		WATSON LABS	<u>0.15MG;0.03MG</u>	<u>A076915</u>	<u>001</u>	Jul 29, 2005

EMOQUETTE

AB		VINTAGE PHARMS LLC	<u>0.15MG;0.03MG</u>	<u>A076675</u>	<u>001</u>	Feb 25, 2011
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ENSKYCE

AB		LUPIN LTD	<u>0.15MG;0.03MG</u>	<u>A201887</u>	<u>001</u>	Mar 07, 2013
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ISIBLOOM

AB		XIROMED	<u>0.15MG;0.03MG</u>	<u>A202789</u>	<u>001</u>	Aug 12, 2015
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KALLIGA

AB		AUROBINDO PHARMA LTD	<u>0.15MG;0.03MG</u>	<u>A207081</u>	<u>001</u>	May 17, 2017
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KARIVA

AB	!	BARR	<u>0.15MG,N/A;0.02MG,0.01MG</u>	<u>A075863</u>	<u>001</u>	Apr 05, 2002
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KIMIDESS

AB		VINTAGE PHARMS	<u>0.15MG,N/A;0.02MG,0.01MG</u>	<u>A076681</u>	<u>001</u>	Apr 30, 2015
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PIMTREA

AB		NOVAST LABS	<u>0.15MG,N/A;0.02MG,0.01MG</u>	<u>A091247</u>	<u>001</u>	Aug 01, 2013
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VELIVET

AB		DURAMED PHARMS BARR	<u>0.1MG,0.125MG,0.15MG;0.025MG,0.025MG,0.025MG</u>	<u>A076455</u>	<u>001</u>	Feb 24, 2004
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VIORELE

AB		GLENMARK GENERICS	<u>0.15MG,N/A;0.02MG,0.01MG</u>	<u>A091346</u>	<u>001</u>	Apr 02, 2012
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VOLNEA

AB		XIROMED	<u>0.15MG,N/A;0.02MG,0.01MG</u>	<u>A202689</u>	<u>001</u>	Sep 09, 2016
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PRESCRIPTION DRUG PRODUCT LIST

DESONIDE

AEROSOL, FOAM;TOPICAL

VERDESO

+! ALMIRALL 0.05% N021978 001 Sep 19, 2006

CREAM;TOPICAL

DESONIDE

AB ACP NIMBLE **0.05%** **A074027 001** Sep 28, 1992

AB CADILA **0.05%** **A210198 001** Nov 20, 2019

AB GLENMARK PHARMS **0.05%** **A209729 001** Jul 24, 2017

AB +! PERRIGO NEW YORK **0.05%** **N017010 001**

AB TARO **0.05%** **A073548 001** Jun 30, 1992

DESOWEN

AB GALDERMA LABS LP **0.05%** **N019048 001** Dec 14, 1984

GEL;TOPICAL

DESONATE

+! LEO PHARMA AS 0.05% N021844 001 Oct 20, 2006

LOTION;TOPICAL

DESONIDE

AB FOUGERA PHARMS **0.05%** **A075860 001** Mar 19, 2002

AB GLENMARK PHARMS **0.05%** **A209494 001** Sep 26, 2017

AB TARO **0.05%** **A202161 001** Oct 31, 2014

AB TELIGENT PHARMA INC **0.05%** **A207855 001** Sep 28, 2017

DESOWEN

AB ! GALDERMA LABS LP **0.05%** **A072354 001** Jan 24, 1992

OINTMENT;TOPICAL

DESONIDE

AB ALEOR **0.05%** **A212473 001** Oct 23, 2019

AB ENCUBE ETHICALS **0.05%** **A210998 001** Jan 30, 2019

AB FOUGERA PHARMS **0.05%** **A075751 001** Mar 12, 2001

AB GLENMARK PHARMS LTD **0.05%** **A209996 001** Sep 15, 2017

AB HI TECH **0.05%** **A208836 001** Mar 27, 2017

AB +! PERRIGO NEW YORK **0.05%** **N017426 001**

AB TARO **0.05%** **A074254 001** Aug 03, 1994

AB TELIGENT PHARMA INC **0.05%** **A212002 001** Mar 12, 2019

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

AB ATLANTIC

AB AKORN **0.05%** **A203787 001** Jan 06, 2017

AB **0.25%** **A203234 001** Jun 12, 2015

AB COSETTE **0.25%** **A209595 001** Mar 04, 2020

AB FOUGERA PHARMS **0.25%** **A078369 001** Jun 29, 2010

AB LUPIN ATLANTIS **0.05%** **A208163 001** Jan 10, 2017

AB **0.25%** **A208164 001** Jan 09, 2017

AB PERRIGO NEW YORK **0.25%** **A076510 001** Jul 01, 2003

AB RISING **0.05%** **A210980 001** Dec 21, 2018

AB **0.25%** **A205594 001** Jul 02, 2018

TOPICORT

AB ! TARO PHARM INDS LTD **0.05%** **A073210 001** Nov 30, 1990

AB ! **0.25%** **A073193 001** Nov 30, 1990

GEL;TOPICAL

DESOXIMETASONE

AB AKORN **0.05%** **A090727 001** Mar 10, 2011

AB PERRIGO NEW YORK **0.05%** **A077552 001** Jan 09, 2006

AB RISING **0.05%** **A204675 001** Aug 12, 2016

TOPICORT

AB ! TARO PHARM INDS LTD **0.05%** **A074904 001** Jul 14, 1998

OINTMENT;TOPICAL

DESOXIMETASONE

AB ACP NIMBLE **0.25%** **A206740 001** Dec 23, 2016

AB ACTAVIS MID **0.25%** **A204965 001** Nov 07, 2016

AB ATLANTIC

AB AKORN **0.25%** **A201005 001** Apr 24, 2014

AB FOUGERA PHARMS **0.25%** **A078657 001** Sep 28, 2012

AB GLENMARK GENERICS **0.25%** **A202838 001** Sep 20, 2013

AB LUPIN ATLANTIS **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

PRESCRIPTION DRUG PRODUCT LIST

DESOXIMETASONE

OINTMENT; TOPICAL

DESOXIMETASONE

<u>AB</u>	PERRIGO ISRAEL	<u>0.25%</u>	<u>A077770</u>	<u>001</u>	Apr 20, 2015
<u>AB</u>	RISING	<u>0.25%</u>	<u>A204272</u>	<u>001</u>	Nov 30, 2016
<u>AB</u>	TELLIGENT PHARMA INC	<u>0.05%</u>	<u>A209973</u>	<u>001</u>	Oct 23, 2018
<u>AB</u>		<u>0.25%</u>	<u>A208101</u>	<u>001</u>	Feb 25, 2016
<u>AB</u>	ZYDUS PHARMS	<u>0.25%</u>	<u>A205206</u>	<u>001</u>	Sep 19, 2017

TOPICORT

<u>AB</u>	+! TARO PHARM INDS LTD	<u>0.05%</u>	<u>N018594</u>	<u>001</u>	Jan 17, 1985
<u>AB</u>	!	<u>0.25%</u>	<u>A074286</u>	<u>001</u>	Jun 07, 1996

SPRAY; TOPICAL

DESOXIMETASONE

<u>AT</u>	LUPIN ATLANTIS	<u>0.25%</u>	<u>A208124</u>	<u>001</u>	Mar 16, 2018
<u>AT</u>	PERRIGO ISRAEL	<u>0.25%</u>	<u>A206441</u>	<u>001</u>	Jan 20, 2017

TOPICORT

<u>AT</u>	+! TARO PHARMS	<u>0.25%</u>	<u>N204141</u>	<u>001</u>	Apr 11, 2013
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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

<u>AB</u>	ACTAVIS LABS FL	<u>EQ 25MG BASE</u>	<u>A204065</u>	<u>001</u>	Jul 29, 2016
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A204065</u>	<u>002</u>	Jul 29, 2016
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A204065</u>	<u>003</u>	Jul 29, 2016
<u>AB</u>	ALEMBIC PHARMS LTD	<u>EQ 25MG BASE</u>	<u>A204003</u>	<u>003</u>	Sep 14, 2018
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A204003</u>	<u>001</u>	Jun 29, 2015
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A204003</u>	<u>002</u>	Jun 29, 2015
<u>AB</u>	CASI PHARMS INC	<u>EQ 50MG BASE</u>	<u>A204028</u>	<u>001</u>	Jun 29, 2015
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A204028</u>	<u>002</u>	Jun 29, 2015
<u>AB</u>	HIKMA	<u>EQ 25MG BASE</u>	<u>A204082</u>	<u>002</u>	Aug 28, 2017
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A204082</u>	<u>001</u>	Feb 16, 2016
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A204083</u>	<u>001</u>	Feb 16, 2016
<u>AB</u>	INTELLIPHARMACEUTIC S	<u>EQ 50MG BASE</u>	<u>A204805</u>	<u>001</u>	May 07, 2019
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A204805</u>	<u>002</u>	May 07, 2019
<u>AB</u>	LUPIN LTD	<u>EQ 50MG BASE</u>	<u>A204172</u>	<u>001</u>	Jun 29, 2015
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A204172</u>	<u>002</u>	Jun 29, 2015
<u>AB</u>	ZYDUS PHARMS	<u>EQ 50MG BASE</u>	<u>A204020</u>	<u>001</u>	Oct 11, 2017
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A204020</u>	<u>002</u>	Oct 11, 2017
	<u>PRISTIO</u>				
<u>AB</u>	+ PF PRISM CV	<u>EQ 25MG BASE</u>	<u>N021992</u>	<u>003</u>	Aug 20, 2014
<u>AB</u>	+!	<u>EQ 50MG BASE</u>	<u>N021992</u>	<u>001</u>	Feb 29, 2008
<u>AB</u>	+!	<u>EQ 100MG BASE</u>	<u>N021992</u>	<u>002</u>	Feb 29, 2008

DEUTETRABENAZINE

TABLET; ORAL

AUSTEDO

	+ TEVA BRANDED PHARM	6MG	N208082	001	Apr 03, 2017
	+	9MG	N208082	002	Apr 03, 2017
	+!	12MG	N208082	003	Apr 03, 2017

DEXAMETHASONE

CONCENTRATE; ORAL

DEXAMETHASONE INTENSOL

	! HIKMA	1MG/ML	A088252	001	Sep 01, 1983
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ELIXIR; ORAL

DEXAMETHASONE

<u>AA</u>	! ANIMA	<u>0.5MG/5ML</u>	<u>A084754</u>	<u>001</u>	
<u>AA</u>	LANNETT CO INC	<u>0.5MG/5ML</u>	<u>A091188</u>	<u>001</u>	May 11, 2011
<u>AA</u>	LYNE	<u>0.5MG/5ML</u>	<u>A090891</u>	<u>001</u>	Jul 12, 2011
<u>AA</u>	WOCKHARDT BIO AG	<u>0.5MG/5ML</u>	<u>A088254</u>	<u>001</u>	Jul 27, 1983

IMPLANT; INTRAVITREAL

OZURDEX

	+! ALLERGAN	0.7MG	N022315	001	Jun 17, 2009
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INSERT; OPHTHALMIC

DEXTENZA

	+! OCULAR THERAPEUTIX	0.4MG	N208742	001	Nov 30, 2018
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PRESCRIPTION DRUG PRODUCT LIST

DEXAMETHASONE

SOLUTION; ORAL

DEXAMETHASONE

! HIKMA

0.5MG/5ML

A088248 001 Sep 01, 1983

SUSPENSION; INTRAOCULAR

DEXYCU KIT

+! EYEPOINT PHARMS

9%

N208912 001 Feb 09, 2018

SUSPENSION/DROPS; OPHTHALMIC

MAXIDEX

+! NOVARTIS

0.1%

N013422 001

TABLET; ORAL

DEXAMETHASONE

AB	ECR	1.5MG	A040700 001	Aug 15, 2008
AB	HIKMA	1.5MG	A084610 001	
AB	LARKEN LABS INC	1.5MG	A201270 001	Jul 17, 2017
BP	FERA PHARMS LLC	0.5MG	A088481 002	Apr 28, 1983
BP		0.75MG	A088481 003	Apr 28, 1983
BP		4MG	A088481 004	Apr 28, 1983
BP		6MG	A088481 001	Nov 28, 1983
BP	HIKMA	0.5MG	A084611 001	
BP		0.75MG	A084613 001	
BP		1MG	A088306 001	Sep 15, 1983
BP		2MG	A087916 001	Aug 26, 1982
BP		4MG	A084612 001	
BP	!	6MG	A088316 001	Sep 15, 1983
BP	XSPIRE PHARMA	1.5MG	A088237 001	Apr 28, 1983
	HEMADY			
	+! DEXCEL PHARMA	20MG	N211379 001	Oct 03, 2019

DEXAMETHASONE SODIUM PHOSPHATE

INJECTABLE; INJECTION

DEXAMETHASONE SODIUM PHOSPHATE

AP	AMNEAL	EQ 4MG PHOSPHATE/ML	A208689 001	Aug 22, 2018
AP	AUROBINDO PHARMA LTD	EQ 4MG PHOSPHATE/ML	A206781 001	Dec 01, 2015
AP	!	EQ 4MG PHOSPHATE/ML	A084916 001	
AP	FRESENIUS KABI USA	EQ 4MG PHOSPHATE/ML	A203129 001	Sep 30, 2015
AP	!	EQ 10MG PHOSPHATE/ML	A040572 001	Apr 22, 2005
AP		EQ 10MG PHOSPHATE/ML	A209192 001	Jul 06, 2018
AP	MYLAN LABS LTD	EQ 4MG PHOSPHATE/ML	A040803 001	Aug 29, 2008
AP		EQ 10MG PHOSPHATE/ML	A040802 001	Aug 29, 2008
AP	SOMERSET	EQ 4MG PHOSPHATE/ML	A207521 001	Jun 08, 2018
AP	SOMERSET THERAPS LLC	EQ 10MG PHOSPHATE/ML	A211036 001	May 10, 2019
AP	WEST-WARD PHARMS INT	EQ 4MG PHOSPHATE/ML	A084282 001	
AP	!	EQ 10MG PHOSPHATE/ML	A087702 001	Sep 07, 1982
	<u>DEXAMETHASONE SODIUM PHOSPHATE PRESERVATIVE FREE</u>			
AP	AMNEAL	EQ 10MG PHOSPHATE/ML	A208690 001	Aug 22, 2018
AP	!	EQ 10MG PHOSPHATE/ML	A040491 001	Apr 11, 2003
AP	SOMERSET THERAPS LLC	EQ 10MG PHOSPHATE/ML	A207442 001	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

AT	+! NOVARTIS	0.1%;EQ 3.5MG BASE/GM;10,000 UNITS/GM	N050065 002	
	<u>NEOMYCIN AND POLYMYXIN B SULFATES AND DEXAMETHASONE</u>			
AT	BAUSCH AND LOMB	0.1%;EQ 3.5MG BASE/GM;10,000 UNITS/GM	A064063 001	Jul 25, 1994
AT	PERRIGO CO TENNESSEE	0.1%;EQ 3.5MG BASE/GM;10,000 UNITS/GM	A062938 001	Jul 31, 1989
	SUSPENSION/DROPS; OPHTHALMIC			
	<u>DEXASPORIN</u>			
AT	BAUSCH AND LOMB	0.1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML	A064135 001	Sep 13, 1995
	<u>MAXITROL</u>			
AT	+! NOVARTIS	0.1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML	N050023 002	
AT	SANDOZ INC	0.1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML	A062341 001	May 22, 1984

PRESCRIPTION DRUG PRODUCT LIST

DEXAMETHASONE; TOBRAMYCIN

OINTMENT;OPHTHALMIC

TOBRADEX

+! NOVARTIS 0.1%;0.3%

N050616 001 Sep 28, 1988

SUSPENSION/DROPS;OPHTHALMIC

TOBRADEX**AB** +! NOVARTIS 0.1%;0.3%**N050592 001** Aug 18, 1988TOBRAMYCIN AND DEXAMETHASONE**AB** BAUSCH AND LOMB 0.1%;0.3%**A064134 001** Oct 27, 1999

TOBRADEX ST

+! EYEVANCE 0.05%;0.3%

N050818 001 Feb 13, 2009

DEXCHLORPHENIRAMINE MALEATE

SYRUP;ORAL

DEXCHLORPHENIRAMINE MALEATE**AA** ! WOCKHARDT BIO AG 2MG/5ML**A088251 001** Mar 23, 1984POLMON**AA** CAPELLON PHARMS LLC 2MG/5ML**A202520 001** Jul 16, 2018DEXLANSOPRAZOLE

CAPSULE, DELAYED RELEASE;ORAL

DEXILANT**AB** +! TAKEDA PHARMS USA 60MG**N022287 002** Jan 30, 2009DEXLANSOPRAZOLE**AB** PAR PHARM INC 60MG**A202294 001** Apr 19, 2017

DEXILANT

+ TAKEDA PHARMS USA 30MG

N022287 001 Jan 30, 2009

DEXMEDETOMIDINE HYDROCHLORIDE

INJECTABLE;INJECTION

DEXMEDETOMIDINE**AP** JIANGSU HENGRUI MED EQ 200MCG BASE/2ML (EQ 100MCG BASE/ML)**A209065 001** Sep 19, 2017DEXMEDETOMIDINE HYDROCHLORIDE**AP** ACCORD HLTHCARE EQ 200MCG BASE/2ML (EQ 100MCG BASE/ML)**A204023 001** Feb 09, 2016**AP** ACTAVIS INC EQ 200MCG BASE/2ML (EQ 100MCG BASE/ML)**A204686 001** Oct 17, 2016**AP** AKORN INC EQ 200MCG BASE/2ML (EQ 100MCG BASE/ML)**A202585 001** Nov 24, 2014**AP** AUROBINDO PHARMA EQ 200MCG BASE/2ML (EQ 100MCG BASE/ML)**A205867 001** Mar 17, 2016**AP** LTD EQ 200MCG BASE/2ML (EQ 100MCG BASE/ML)**AP** BAXTER HLTHCARE EQ 200MCG BASE/50ML (EQ 4MCG BASE/ML)**A208532 001** Aug 21, 2018**AP** CORP EQ 400MCG BASE/100ML (EQ 4MCG BASE/ML)**A208532 002** Aug 21, 2018**AP** FRESENIUS KABI USA EQ 80MCG BASE/20ML (EQ 4MCG BASE/ML)**A208129 001** Nov 29, 2018**AP** EQ 200MCG BASE/2ML (EQ 100MCG BASE/ML)**A201072 001** Sep 18, 2015**AP** EQ 200MCG BASE/50ML (EQ 4MCG BASE/ML)**A208129 002** Nov 29, 2018**AP** EQ 400MCG BASE/100ML (EQ 4MCG BASE/ML)**A208129 003** Nov 29, 2018**AP** HIKMA EQ 200MCG BASE/50ML (EQ 4MCG BASE/ML)**A206407 001** Jan 30, 2020**AP** EQ 400MCG BASE/100ML (EQ 4MCG BASE/ML)**A206407 002** Jan 30, 2020**AP** MYLAN INSTITUTIONAL EQ 200MCG BASE/2ML (EQ 100MCG BASE/ML)**A202881 001** Aug 18, 2014**AP** PAR STERILE EQ 200MCG BASE/2ML (EQ 100MCG BASE/ML)**A203972 001** Aug 18, 2014**AP** PRODUCTS EQ 200MCG BASE/2ML (EQ 100MCG BASE/ML)**AP** SANDOZ INC EQ 200MCG BASE/2ML (EQ 100MCG BASE/ML)**A091465 001** Jun 14, 2016**AP** SLAYBACK PHARMA LLC EQ 200MCG BASE/50ML (EQ 4MCG BASE/ML)**A212791 001** Dec 04, 2019**AP** EQ 400MCG BASE/100ML (EQ 4MCG BASE/ML)**A212791 002** Dec 04, 2019**AP** SUN PHARM INDS INC EQ 200MCG BASE/2ML (EQ 100MCG BASE/ML)**A202126 001** Aug 20, 2015**AP** TEVA PHARMS USA EQ 200MCG BASE/2ML (EQ 100MCG BASE/ML)**A205272 001** Nov 28, 2017**AP** WEST-WARD PHARMS EQ 200MCG BASE/2ML (EQ 100MCG BASE/ML)**A205046 001** Apr 26, 2017**AP** INT EQ 200MCG BASE/2ML (EQ 100MCG BASE/ML)**AP** ZYDUS PHARMS EQ 200MCG BASE/2ML (EQ 100MCG BASE/ML)**A206798 001** Feb 27, 2018PRECEDEX**AP** +! HOSPIRA EQ 80MCG BASE/20ML (EQ 4MCG BASE/ML)**N021038 004** Nov 14, 2014**AP** +! EQ 200MCG BASE/2ML (EQ 100MCG BASE/ML)**N021038 001** Dec 17, 1999**AP** +! EQ 200MCG BASE/50ML (EQ 4MCG BASE/ML)**N021038 002** Mar 13, 2013**AP** +! EQ 400MCG BASE/100ML (EQ 4MCG BASE/ML)**N021038 003** Mar 13, 2013

SOLUTION;INTRAVENOUS

DEXMEDETOMIDINE HYDROCHLORIDE

+! HQ SPCLT PHARMA EQ 1MG BASE/10ML (EQ 100MCG BASE/ML)

N206628 002 Oct 21, 2015

+ EQ 200MCG BASE/50ML (EQ 4MCG BASE/ML)

N206628 003 Jun 22, 2018

+ EQ 400MCG BASE/4ML (EQ 100MCG BASE/ML)

N206628 001 Oct 21, 2015

+ EQ 400MCG BASE/100ML (EQ 4MCG BASE/ML)

N206628 004 Jun 22, 2018

PRESCRIPTION DRUG PRODUCT LIST

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>	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>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>	MYLAN	<u>5MG</u>	<u>A204266 001</u>	Aug 25, 2015
<u>AB</u>		<u>10MG</u>	<u>A204266 002</u>	Aug 25, 2015
<u>AB</u>		<u>15MG</u>	<u>A204266 003</u>	Aug 25, 2015
<u>AB</u>		<u>20MG</u>	<u>A204266 004</u>	Dec 21, 2015
<u>AB</u>		<u>30MG</u>	<u>A202580 001</u>	Aug 28, 2013
<u>AB</u>		<u>40MG</u>	<u>A204266 007</u>	Aug 25, 2015
<u>AB</u>	PAR PHARM INC	<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>	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
<u>FOCALIN XR</u>				
<u>AB</u>	+ NOVARTIS	<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
<u>AB</u>		<u>10MG</u>	<u>A204534 003</u>	Dec 04, 2015
<u>AB</u>	RHODES PHARMS	<u>2.5MG</u>	<u>A208756 001</u>	Nov 20, 2017
<u>AB</u>		<u>5MG</u>	<u>A208756 002</u>	Nov 20, 2017
<u>AB</u>		<u>10MG</u>	<u>A208756 003</u>	Nov 20, 2017
<u>AB</u>	SUN PHARM INDUSTRIES	<u>2.5MG</u>	<u>A201231 001</u>	Sep 24, 2015
<u>AB</u>		<u>5MG</u>	<u>A201231 002</u>	Sep 24, 2015
<u>AB</u>		<u>10MG</u>	<u>A201231 003</u>	Sep 24, 2015
<u>AB</u>	TRIS PHARMA INC	<u>2.5MG</u>	<u>A207901 001</u>	Aug 26, 2016

PRESCRIPTION DRUG PRODUCT LIST

DEXMETHYLPHENIDATE HYDROCHLORIDE

TABLET; ORAL

DEXMETHYLPHENIDATE HYDROCHLORIDE

<u>AB</u>		<u>5MG</u>	<u>A207901 002</u>	Aug 26, 2016
<u>AB</u>		<u>10MG</u>	<u>A207901 003</u>	Aug 26, 2016
<u>FOCALIN</u>				
<u>AB</u>	+	NOVARTIS	<u>2.5MG</u>	<u>N021278 001</u> Nov 13, 2001
<u>AB</u>	+		<u>5MG</u>	<u>N021278 002</u> Nov 13, 2001
<u>AB</u>	+	!	<u>10MG</u>	<u>N021278 003</u> Nov 13, 2001

DEXRAZOXANE HYDROCHLORIDE

INJECTABLE; INJECTION

DEXRAZOXANE HYDROCHLORIDE

<u>AP</u>		GLAND PHARMA LTD	<u>EQ 250MG BASE/VIAL</u>	<u>A207321 002</u> Dec 16, 2019
<u>AP</u>			<u>EQ 500MG BASE/VIAL</u>	<u>A207321 001</u> Nov 28, 2016
<u>AP</u>		MYLAN INSTITUTIONAL	<u>EQ 250MG BASE/VIAL</u>	<u>A200752 001</u> Oct 19, 2011
<u>AP</u>			<u>EQ 500MG BASE/VIAL</u>	<u>A200752 002</u> Oct 19, 2011
<u>AP</u>		WEST-WARD PHARMS INT	<u>EQ 250MG BASE/VIAL</u>	<u>A076068 001</u> Sep 28, 2004
<u>AP</u>			<u>EQ 500MG BASE/VIAL</u>	<u>A076068 002</u> Sep 28, 2004
<u>ZINECARD</u>				
<u>AP</u>	+	PHARMACIA AND UPJOHN	<u>EQ 250MG BASE/VIAL</u>	<u>N020212 001</u> May 26, 1995
<u>AP</u>	+	!	<u>EQ 500MG BASE/VIAL</u>	<u>N020212 002</u> May 26, 1995
TOTECT				
	+	CLINIGEN	<u>EQ 500MG BASE/VIAL</u>	<u>N022025 001</u> Sep 06, 2007

DEXTROAMPHETAMINE SULFATE

CAPSULE, EXTENDED RELEASE; ORAL

DEXEDRINE

<u>AB</u>	+	IMPAX LABS INC	<u>5MG</u>	<u>N017078 001</u>
<u>AB</u>	+		<u>10MG</u>	<u>N017078 002</u>
<u>AB</u>	+	!	<u>15MG</u>	<u>N017078 003</u>

DEXTROAMPHETAMINE SULFATE

<u>AB</u>		ACTAVIS ELIZABETH	<u>5MG</u>	<u>A203901 001</u> Nov 30, 2012
<u>AB</u>			<u>10MG</u>	<u>A203901 002</u> Nov 30, 2012
<u>AB</u>			<u>15MG</u>	<u>A203901 003</u> Nov 30, 2012
<u>AB</u>		MAYNE PHARMA	<u>5MG</u>	<u>A076137 001</u> Jan 18, 2002
<u>AB</u>			<u>10MG</u>	<u>A076137 002</u> Jan 18, 2002
<u>AB</u>			<u>15MG</u>	<u>A076137 003</u> Jan 18, 2002
<u>AB</u>		NESHER PHARMS	<u>5MG</u>	<u>A209111 001</u> Jun 27, 2017
<u>AB</u>			<u>10MG</u>	<u>A209111 002</u> Jun 27, 2017
<u>AB</u>			<u>15MG</u>	<u>A209111 003</u> Jun 27, 2017
<u>AB</u>		SPECGX LLC	<u>5MG</u>	<u>A076353 001</u> May 06, 2003
<u>AB</u>			<u>10MG</u>	<u>A076353 002</u> May 06, 2003
<u>AB</u>			<u>15MG</u>	<u>A076353 003</u> May 06, 2003
<u>AB</u>		VINTAGE PHARMS	<u>5MG</u>	<u>A205673 001</u> Oct 31, 2017
<u>AB</u>			<u>10MG</u>	<u>A205673 002</u> Oct 31, 2017
<u>AB</u>			<u>15MG</u>	<u>A205673 003</u> Oct 31, 2017

SOLUTION; ORAL

DEXTROAMPHETAMINE SULFATE

<u>AA</u>	!	OUTLOOK PHARMS	<u>5MG/5ML</u>	<u>A040776 001</u> Jan 29, 2008
<u>AA</u>		TRIS PHARMA INC	<u>5MG/5ML</u>	<u>A203644 001</u> May 29, 2013

TABLET; ORAL

DEXTROAMPHETAMINE SULFATE

<u>AA</u>		ARBOR PHARMS LLC	<u>5MG</u>	<u>A090533 002</u> Oct 25, 2011
<u>AA</u>			<u>10MG</u>	<u>A090533 004</u> Oct 25, 2011
<u>AA</u>		AUROLIFE PHARMA LLC	<u>5MG</u>	<u>A202893 001</u> Jul 31, 2013
<u>AA</u>			<u>10MG</u>	<u>A202893 002</u> Jul 31, 2013
<u>AA</u>		AVANTHI INC	<u>5MG</u>	<u>A203548 001</u> Nov 23, 2015
<u>AA</u>			<u>10MG</u>	<u>A203548 002</u> Nov 23, 2015
<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>		NESHER PHARMS	<u>5MG</u>	<u>A206588 001</u> Mar 28, 2016
<u>AA</u>			<u>10MG</u>	<u>A206588 002</u> Mar 28, 2016
<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
		ARBOR PHARMS LLC	2.5MG	A090533 001 Oct 25, 2011
			7.5MG	A090533 003 Oct 25, 2011
			15MG	A090533 005 Oct 25, 2011

PRESCRIPTION DRUG PRODUCT LISTDEXTROAMPHETAMINE SULFATE

TABLET; ORAL

DEXTROAMPHETAMINE SULFATE

20MG

A090533 006 Oct 25, 2011

30MG

A090533 007 Oct 25, 2011

DEXTROMETHORPHAN HYDROBROMIDE; PROMETHAZINE HYDROCHLORIDE

SYRUP; ORAL

PROMETHAZINE HYDROCHLORIDE AND DEXTROMETHORPHAN HYDROBROMIDE**AA** HI TECH PHARMA 15MG/5ML; 6.25MG/5ML **A040027 001** Jul 31, 1996PROMETHAZINE W/ DEXTROMETHORPHAN**AA** WOCKHARDT BIO AG 15MG/5ML; 6.25MG/5ML **A088864 001** Jan 04, 1985DEXTROMETHORPHAN HYDROBROMIDE; QUINIDINE SULFATE

CAPSULE; ORAL

DEXTROMETHORPHAN HYDROBROMIDE AND QUINIDINE SULFATE**AB** ACTAVIS ELIZABETH 20MG; 10MG **A202934 001** Oct 10, 2017NUDEXTA**AB** +! AVANIR PHARMS 20MG; 10MG **N021879 001** Oct 29, 2010DEXTROSE

INJECTABLE; INJECTION

DEXTROSE 10% IN PLASTIC CONTAINER**AP** +! B BRAUN 10GM/100ML **N019626 004** Feb 02, 1988**AP** +! BAXTER HLTHCARE 10GM/100ML **N016694 001****AP** FRESENIUS KABI USA 10GM/100ML **A209448 001** Jul 16, 2018**AP** +! ICU MEDICAL INC 10GM/100ML **N018080 001**DEXTROSE 5% IN PLASTIC CONTAINER**AP** +! B BRAUN 50MG/ML **N016730 002****AP** +! 5GM/100ML **N016730 001****AP** +! 5GM/100ML **N019626 002** Feb 02, 1988**AP** +! BAXTER HLTHCARE 50MG/ML **N016673 003** Oct 30, 1985**AP** +! 50MG/ML **N020179 002** Dec 07, 1992**AP** +! 5GM/100ML **N016673 001****AP** +! 5GM/100ML **N020179 001** Dec 07, 1992**AP** FRESENIUS KABI USA 50MG/ML **A207449 001** Oct 21, 2016**AP** +! HOSPIRA 5GM/100ML **N019466 001** Jul 15, 1985**AP** +! 5GM/100ML **N019479 001** Sep 17, 1985**AP** +! ICU MEDICAL INC 50MG/ML **N016367 002**DEXTROSE 50% IN PLASTIC CONTAINER**AP** +! BAXTER HLTHCARE 50GM/100ML **N020047 001** Jul 02, 1991**AP** +! ICU MEDICAL INC 50GM/100ML **N018563 001** Mar 23, 1982DEXTROSE 70% IN PLASTIC CONTAINER**AP** +! B BRAUN 70GM/100ML **N019626 005** Feb 18, 2015**AP** +! BAXTER HLTHCARE 70GM/100ML **N020047 003** Jul 02, 1991**AP** +! ICU MEDICAL INC 70GM/100ML **N018561 001** Mar 23, 1982**AP** +! 70GM/100ML **N019893 001** Dec 26, 1989

DEXTROSE 20% IN PLASTIC CONTAINER

+! ICU MEDICAL INC 20GM/100ML N018564 001 Mar 23, 1982

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 50%

+ HOSPIRA 500MG/ML N019445 003 Sep 03, 2014

DEXTROSE 50% IN PLASTIC CONTAINER

+ HOSPIRA 500MG/ML N019445 001 Jun 03, 1986

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

PRESCRIPTION DRUG PRODUCT LIST

<u>DEXTROSE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; POTASSIUM PHOSPHATE, MONOBASIC; SODIUM CHLORIDE; SODIUM LACTATE</u>			
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
<u>DEXTROSE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; POTASSIUM PHOSPHATE, MONOBASIC; SODIUM LACTATE; SODIUM PHOSPHATE, MONOBASIC ANHYDROUS</u>			
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
<u>DEXTROSE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE; SODIUM GLUCONATE</u>			
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
<u>DEXTROSE; POTASSIUM CHLORIDE</u>			
INJECTABLE; INJECTION			
<u>DEXTROSE 5% AND POTASSIUM CHLORIDE 0.15% IN PLASTIC CONTAINER</u>			
AP	+ BAXTER HLTHCARE	5GM/100ML; 150MG/100ML	N017634 001
<u>DEXTROSE 5% AND POTASSIUM CHLORIDE 0.224% IN PLASTIC CONTAINER</u>			
AP	+ BAXTER HLTHCARE	5GM/100ML; 224MG/100ML	N017634 003
<u>DEXTROSE 5% AND POTASSIUM CHLORIDE 0.3% IN PLASTIC CONTAINER</u>			
AP	+ BAXTER HLTHCARE	5GM/100ML; 300MG/100ML	N017634 002
<u>POTASSIUM CHLORIDE 0.15% IN DEXTROSE 5% IN PLASTIC CONTAINER</u>			
AP	B BRAUN	5GM/100ML; 150MG/100ML	N019699 004 Sep 29, 1989
<u>POTASSIUM CHLORIDE 0.3% IN DEXTROSE 5% IN PLASTIC CONTAINER</u>			
AP	B BRAUN	5GM/100ML; 300MG/100ML	N019699 006 Sep 29, 1989
DEXTROSE 5% AND POTASSIUM CHLORIDE 0.075% IN PLASTIC CONTAINER			
	+ BAXTER HLTHCARE	5GM/100ML; 75MG/100ML	N017634 004
POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% IN PLASTIC CONTAINER			
	ICU MEDICAL INC	5GM/100ML; 149MG/100ML	N018371 001
<u>DEXTROSE; POTASSIUM CHLORIDE; SODIUM CHLORIDE</u>			
INJECTABLE; INJECTION			
<u>DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 10MEQ</u>			
AP	BAXTER HLTHCARE	5GM/100ML; 75MG/100ML; 200MG/100ML	N018037 006 Apr 13, 1982
AP		5GM/100ML; 150MG/100ML; 200MG/100ML	N018037 007 Apr 13, 1982
<u>DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 15MEQ (K)</u>			
AP	BAXTER HLTHCARE	5GM/100ML; 224MG/100ML; 200MG/100ML	N018037 004
<u>DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 20MEQ</u>			
AP	BAXTER HLTHCARE	5GM/100ML; 150MG/100ML; 200MG/100ML	N018037 008 Apr 13, 1982
<u>DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 20MEQ (K)</u>			
AP	BAXTER HLTHCARE	5GM/100ML; 300MG/100ML; 200MG/100ML	N018037 001
<u>DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 30MEQ</u>			
AP	BAXTER HLTHCARE	5GM/100ML; 224MG/100ML; 200MG/100ML	N018037 005 Apr 13, 1982
<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

PRESCRIPTION DRUG PRODUCT LIST

DEXTROSE; POTASSIUM CHLORIDE; SODIUM CHLORIDE

INJECTABLE; INJECTION

		<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 0.45% 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 0.45% 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% 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
		<u>POTASSIUM CHLORIDE 30MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER</u>		
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	
		<u>POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER</u>		
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	
		<u>POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER</u>		
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

PRESCRIPTION DRUG PRODUCT LISTDEXTROSE; POTASSIUM CHLORIDE; SODIUM CHLORIDE

INJECTABLE; INJECTION

POTASSIUM CHLORIDE 0.11% IN B BRAUN	DEXTROSE 10% AND SODIUM CHLORIDE 0.45% IN 10GM/100ML;110MG/100ML;450MG/100ML	IN PLASTIC CONTAINER N019630 039 Feb 17, 1988
POTASSIUM CHLORIDE 0.11% IN B BRAUN	DEXTROSE 10% AND SODIUM CHLORIDE 0.9% IN 10GM/100ML;110MG/100ML;900MG/100ML	IN PLASTIC CONTAINER N019630 045 Feb 17, 1988
POTASSIUM CHLORIDE 0.11% IN B BRAUN	DEXTROSE 3.3% AND SODIUM CHLORIDE 0.3% IN 3.3GM/100ML;110MG/100ML;300MG/100ML	IN PLASTIC CONTAINER N019630 050 May 07, 1992
POTASSIUM CHLORIDE 0.11% IN B BRAUN	DEXTROSE 5% AND SODIUM CHLORIDE 0.11% IN 5GM/100ML;110MG/100ML;110MG/100ML	IN PLASTIC CONTAINER N019630 003 Feb 17, 1988
POTASSIUM CHLORIDE 0.11% IN B BRAUN	DEXTROSE 5% AND SODIUM CHLORIDE 0.2% IN 5GM/100ML;110MG/100ML;200MG/100ML	IN PLASTIC CONTAINER N019630 009 Feb 17, 1988
POTASSIUM CHLORIDE 0.11% IN B BRAUN	DEXTROSE 5% AND SODIUM CHLORIDE 0.33% IN 5GM/100ML;110MG/100ML;330MG/100ML	IN PLASTIC CONTAINER N019630 015 Feb 17, 1988
POTASSIUM CHLORIDE 0.11% IN B BRAUN	DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN 5GM/100ML;110MG/100ML;450MG/100ML	IN PLASTIC CONTAINER N019630 021 Feb 17, 1988
POTASSIUM CHLORIDE 0.11% IN B BRAUN	DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN 5GM/100ML;110MG/100ML;900MG/100ML	IN PLASTIC CONTAINER N019630 027 Feb 17, 1988
POTASSIUM CHLORIDE 0.15% IN B BRAUN	DEXTROSE 10% AND SODIUM CHLORIDE 0.2% IN 10GM/100ML;150MG/100ML;200MG/100ML	IN PLASTIC CONTAINER N019630 034 Feb 17, 1988
POTASSIUM CHLORIDE 0.15% IN B BRAUN	DEXTROSE 10% AND SODIUM CHLORIDE 0.45% IN 10GM/100ML;150MG/100ML;450MG/100ML	IN PLASTIC CONTAINER N019630 040 Feb 17, 1988
POTASSIUM CHLORIDE 0.15% IN B BRAUN	DEXTROSE 10% AND SODIUM CHLORIDE 0.9% IN 10GM/100ML;150MG/100ML;900MG/100ML	IN PLASTIC CONTAINER N019630 046 Feb 17, 1988
POTASSIUM CHLORIDE 0.15% IN B BRAUN	DEXTROSE 3.3% AND SODIUM CHLORIDE 0.3% IN 3.3GM/100ML;150MG/100ML;300MG/100ML	IN PLASTIC CONTAINER N019630 051 May 07, 1992
POTASSIUM CHLORIDE 0.15% IN B BRAUN	DEXTROSE 5% AND SODIUM CHLORIDE 0.11% IN 5GM/100ML;150MG/100ML;110MG/100ML	IN PLASTIC CONTAINER N019630 004 Feb 17, 1988
POTASSIUM CHLORIDE 0.22% IN B BRAUN	DEXTROSE 10% AND SODIUM CHLORIDE 0.2% IN 10GM/100ML;220MG/100ML;200MG/100ML	IN PLASTIC CONTAINER N019630 035 Feb 17, 1988
POTASSIUM CHLORIDE 0.22% IN B BRAUN	DEXTROSE 10% AND SODIUM CHLORIDE 0.45% IN 10GM/100ML;220MG/100ML;450MG/100ML	IN PLASTIC CONTAINER N019630 041 Feb 17, 1988
POTASSIUM CHLORIDE 0.22% IN B BRAUN	DEXTROSE 10% AND SODIUM CHLORIDE 0.9% IN 10GM/100ML;220MG/100ML;900MG/100ML	IN PLASTIC CONTAINER N019630 047 Feb 17, 1988
POTASSIUM CHLORIDE 0.22% IN B BRAUN	DEXTROSE 3.3% AND SODIUM CHLORIDE 0.3% IN 3.3GM/100ML;220MG/100ML;300MG/100ML	IN PLASTIC CONTAINER N019630 052 May 07, 1992
POTASSIUM CHLORIDE 0.22% IN B BRAUN	DEXTROSE 5% AND SODIUM CHLORIDE 0.11% IN 5GM/100ML;220MG/100ML;110MG/100ML	IN PLASTIC CONTAINER N019630 005 Feb 17, 1988
POTASSIUM CHLORIDE 0.22% IN B BRAUN	DEXTROSE 5% AND SODIUM CHLORIDE 0.2% IN 5GM/100ML;220MG/100ML;200MG/100ML	IN PLASTIC CONTAINER N019630 011 Feb 17, 1988
POTASSIUM CHLORIDE 0.22% IN B BRAUN	DEXTROSE 5% AND SODIUM CHLORIDE 0.33% IN 5GM/100ML;220MG/100ML;330MG/100ML	IN PLASTIC CONTAINER N019630 017 Feb 17, 1988
POTASSIUM CHLORIDE 0.22% IN B BRAUN	DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN 5GM/100ML;220MG/100ML;450MG/100ML	IN PLASTIC CONTAINER N019630 023 Feb 17, 1988
POTASSIUM CHLORIDE 0.22% IN B BRAUN	DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN 5GM/100ML;220MG/100ML;900MG/100ML	IN PLASTIC CONTAINER N019630 029 Feb 17, 1988
POTASSIUM CHLORIDE 0.3% IN B BRAUN	DEXTROSE 10% AND SODIUM CHLORIDE 0.2% IN 10GM/100ML;300MG/100ML;200MG/100ML	IN PLASTIC CONTAINER N019630 036 Feb 17, 1988
POTASSIUM CHLORIDE 0.3% IN B BRAUN	DEXTROSE 10% AND SODIUM CHLORIDE 0.45% IN 10GM/100ML;300MG/100ML;450MG/100ML	IN PLASTIC CONTAINER N019630 042 Feb 17, 1988
POTASSIUM CHLORIDE 0.3% IN B BRAUN	DEXTROSE 10% AND SODIUM CHLORIDE 0.9% IN 10GM/100ML;300MG/100ML;900MG/100ML	IN PLASTIC CONTAINER N019630 048 Feb 17, 1988
POTASSIUM CHLORIDE 0.3% IN B BRAUN	DEXTROSE 3.3% AND SODIUM CHLORIDE 0.3% IN 3.3GM/100ML;300MG/100ML;300MG/100ML	IN PLASTIC CONTAINER N019630 053 May 07, 1992
POTASSIUM CHLORIDE 0.3% IN B BRAUN	DEXTROSE 5% AND SODIUM CHLORIDE 0.11% IN 5GM/100ML;300MG/100ML;110MG/100ML	IN PLASTIC CONTAINER N019630 006 Feb 17, 1988
POTASSIUM CHLORIDE 10MEQ IN BAXTER HLTHCARE	DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN 5GM/100ML;75MG/100ML;900MG/100ML	IN PLASTIC CONTAINER N019308 004 Apr 05, 1985
POTASSIUM CHLORIDE 20MEQ IN + ICU MEDICAL INC	DEXTROSE 5% AND SODIUM CHLORIDE 0.225% IN 5GM/100ML;149MG/100ML;225MG/100ML	IN PLASTIC CONTAINER N018365 001
POTASSIUM CHLORIDE 20MEQ IN BAXTER HLTHCARE	DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN 5GM/100ML;300MG/100ML;900MG/100ML	IN PLASTIC CONTAINER N019308 003 Apr 05, 1985
POTASSIUM CHLORIDE 30MEQ IN BAXTER HLTHCARE	DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN 5GM/100ML;224MG/100ML;900MG/100ML	IN PLASTIC CONTAINER N019308 006 Apr 05, 1985
POTASSIUM CHLORIDE 5MEQ IN BAXTER HLTHCARE	DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN 5GM/100ML;150MG/100ML;450MG/100ML	IN PLASTIC CONTAINER N018008 004
POTASSIUM CHLORIDE 5MEQ IN BAXTER HLTHCARE	DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN 5GM/100ML;150MG/100ML;900MG/100ML	IN PLASTIC CONTAINER N019308 001 Apr 05, 1985

PRESCRIPTION DRUG PRODUCT LIST

DEXTROSE; SODIUM CHLORIDE

INJECTABLE; INJECTION

DEXTROSE 2.5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER

AP	B BRAUN	<u>2.5GM/100ML;450MG/100ML</u>	N019631 004	Feb 24, 1988
AP	+! BAXTER HLTHCARE	<u>2.5GM/100ML;450MG/100ML</u>	N016697 001	
AP	FRESENIUS KABI USA	<u>2.5GM/100ML;450MG/100ML</u>	A211190 001	Dec 20, 2019
<u>DEXTROSE 5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER</u>				
AP	B BRAUN	<u>5GM/100ML;200MG/100ML</u>	N019631 007	Feb 24, 1988
<u>DEXTROSE 5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER</u>				
AP	B BRAUN	<u>5GM/100ML;330MG/100ML</u>	N019631 008	Feb 24, 1988
<u>DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER</u>				
AP	B BRAUN	<u>5GM/100ML;450MG/100ML</u>	N019631 009	Feb 24, 1988
AP	+ ICU MEDICAL INC	<u>5GM/100ML;450MG/100ML</u>	N017607 001	
<u>DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER</u>				
AP	B BRAUN	<u>5GM/100ML;900MG/100ML</u>	N019631 010	Feb 24, 1988
AP	+! ICU MEDICAL INC	<u>5GM/100ML;900MG/100ML</u>	N017585 001	
<u>DEXTROSE 5% IN SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER</u>				
AP	BAXTER HLTHCARE	<u>5GM/100ML;200MG/100ML</u>	N016689 001	
<u>DEXTROSE 5% IN SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER</u>				
AP	BAXTER HLTHCARE	<u>5GM/100ML;330MG/100ML</u>	N016687 001	
<u>DEXTROSE 5% IN SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER</u>				
AP	BAXTER HLTHCARE	<u>5GM/100ML;450MG/100ML</u>	N016683 001	
<u>DEXTROSE 5% IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER</u>				
AP	BAXTER HLTHCARE	<u>5GM/100ML;900MG/100ML</u>	N016678 001	
<u>DEXTROSE 10% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER</u>				
	B BRAUN	10GM/100ML;110MG/100ML	N019631 011	Feb 24, 1988
<u>DEXTROSE 10% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER</u>				
	B BRAUN	10GM/100ML;200MG/100ML	N019631 012	Feb 24, 1988
<u>DEXTROSE 10% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER</u>				
	B BRAUN	10GM/100ML;330MG/100ML	N019631 013	Feb 24, 1988
<u>DEXTROSE 10% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER</u>				
	B BRAUN	10GM/100ML;450MG/100ML	N019631 014	Feb 24, 1988
<u>DEXTROSE 10% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER</u>				
	B BRAUN	10GM/100ML;900MG/100ML	N019631 015	Feb 24, 1988
<u>DEXTROSE 2.5% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER</u>				
	B BRAUN	2.5GM/100ML;110MG/100ML	N019631 001	Feb 24, 1988
<u>DEXTROSE 2.5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER</u>				
	B BRAUN	2.5GM/100ML;200MG/100ML	N019631 002	Feb 24, 1988
<u>DEXTROSE 2.5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER</u>				
	B BRAUN	2.5GM/100ML;330MG/100ML	N019631 003	Feb 24, 1988
<u>DEXTROSE 2.5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER</u>				
	B BRAUN	2.5GM/100ML;900MG/100ML	N019631 005	Feb 24, 1988
<u>DEXTROSE 3.3% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER</u>				
	B BRAUN	3.3GM/100ML;300MG/100ML	N019631 016	Jan 19, 1990
<u>DEXTROSE 5% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER</u>				
	B BRAUN	5GM/100ML;110MG/100ML	N019631 006	Feb 24, 1988
<u>DEXTROSE 5% AND SODIUM CHLORIDE 0.225% IN PLASTIC CONTAINER</u>				
	+! ICU MEDICAL INC	5GM/100ML;225MG/100ML	N017606 001	
<u>DEXTROSE 5% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER</u>				
	+! ICU MEDICAL INC	5GM/100ML;300MG/100ML	N017799 001	

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

GASTROGRAFIN**AA** +! BRACCO **66%;10%** **N011245 003**MD-GASTROVIEW**AA** LIEBEL-FLARSHEIM **66%;10%** **A087388 001**DIAZEPAM

CONCENTRATE; ORAL

DIAZEPAM**AA** LANNETT CO INC **5MG/ML** **A204433 001** Apr 14, 2014DIAZEPAM INTENSOL**AA** ! HIKMA **5MG/ML** **A071415 001** Apr 03, 1987

PRESCRIPTION DRUG PRODUCT LIST

DIAZEPAM

GEL; RECTAL

DIASTAT

+ BAUSCH 2.5MG/0.5ML (5MG/ML) N020648 001 Jul 29, 1997

DIASTAT ACUDIAL

+ BAUSCH 10MG/2ML (5MG/ML) N020648 007 Sep 15, 2005

+! 20MG/4ML (5MG/ML) N020648 006 Sep 15, 2005

INJECTABLE; INJECTION

DIAZEPAMAP BELOTECA INC 10MG/2ML (5MG/ML) A210363 001 Mar 18, 2019AP 50MG/10ML (5MG/ML) A211998 001 Dec 26, 2019AP ! HOSPIRA 10MG/2ML (5MG/ML) A072079 001 Dec 20, 1988AP ! 50MG/10ML (5MG/ML) A071583 001 Oct 13, 1987

SOLUTION; ORAL

DIAZEPAMAA ! HIKMA 5MG/5ML A070928 001 Apr 03, 1987AA LANNETT CO INC 5MG/5ML A206477 001 Jun 24, 2016

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

DIAZEPAMAB BARR 2MG A070152 001 Nov 01, 1985AB 10MG A070154 001 Nov 01, 1985AB IVAX SUB TEVA 2MG A071307 001 Dec 10, 1986

PHARMS

AB 5MG A071321 001 Dec 10, 1986AB 10MG A071322 001 Dec 10, 1986AB MAYNE PHARMA 2MG A071134 001 Feb 03, 1987AB 5MG A071135 001 Feb 03, 1987AB 10MG A071136 001 Feb 03, 1987AB MYLAN 2MG A070325 002 Sep 04, 1985AB 5MG A070325 003 Sep 04, 1985AB 10MG A070325 001 Sep 04, 1985AB VINTAGE PHARMS 2MG A077749 001 Mar 31, 2006AB 5MG A077749 002 Mar 31, 2006AB 10MG A077749 003 Mar 31, 2006VALIUMAB + ROCHE 2MG N013263 002AB + 5MG N013263 004AB +! 10MG N013263 006DIAZOXIDE

SUSPENSION; ORAL

DIAZOXIDEAB E5 PHARMA INC 50MG/ML A211050 001 Dec 20, 2019PROGLYCEMAB +! TEVA BRANDED PHARM 50MG/ML N017453 001DICHLORPHENAMIDE

TABLET; ORAL

KEVEYIS

+! STRONGBRIDGE US 50MG N011366 002 Aug 07, 2015

DICLOFENAC

CAPSULE; ORAL

ZORVOLEX

+ ZYLA 18MG N204592 001 Oct 18, 2013

+! 35MG N204592 002 Oct 18, 2013

DICLOFENAC EPOLAMINE

SYSTEM; TOPICAL

FLECTOR

+! INST BIOCHEM 1.3% N021234 001 Jan 31, 2007

DICLOFENAC POTASSIUM

CAPSULE; ORAL

DICLOFENAC POTASSIUMAB BIONPHARMA INC 25MG A204648 001 Feb 23, 2016AB STRIDES PHARMA 25MG A210078 001 Dec 03, 2019ZIPSORAB +! ASSERTIO 25MG N022202 001 Jun 16, 2009

PRESCRIPTION DRUG PRODUCT LIST

DICLOFENAC POTASSIUM

FOR SOLUTION;ORAL

CAMBIA

AB	+ !	ASSERTIO	50MG	N022165	001	Jun 17, 2009
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DICLOFENAC POTASSIUM

AB		PAR FORM	50MG	A202964	001	May 02, 2016
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TABLET;ORAL

DICLOFENAC POTASSIUM

AB		AMICI	50MG	A076561	001	Mar 18, 2004
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AB		CASI PHARMS INC	50MG	A075229	001	Nov 20, 1998
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AB	!	MYLAN	50MG	A075463	001	Jul 26, 1999
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AB		TEVA	50MG	A075219	001	Aug 06, 1998
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DICLOFENAC SODIUM

GEL;TOPICAL

DICLOFENAC SODIUM

AB		ACTAVIS MID	3%	A206493	001	Dec 02, 2015
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ATLANTIC

AB		AMNEAL PHARMS	1%	A208077	001	Mar 18, 2016
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AB		CIPLA	1%	A209903	001	Aug 03, 2018
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AB		ENCUBE	1%	A210986	001	Jan 27, 2020
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AB		GLENMARK PHARMS LTD	3%	A208301	001	Sep 13, 2016
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AB		HI TECH	1%	A209484	001	Nov 21, 2018
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AB		MYLAN	1%	A204306	001	May 06, 2019
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AB		PERRIGO UK FINCO	1%	A211253	001	May 16, 2019
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AB			3%	A210893	001	Jul 27, 2018
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AB		TARO	3%	A206298	001	Apr 28, 2016
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AB		TOLMAR	3%	A200936	001	Oct 28, 2013
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SOLARAZE

AB	+ !	FOUGERA PHARMS	3%	N021005	001	Oct 16, 2000
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VOLTAREN

AB	+ !	GLAXOSMITHKLINE	1%	N022122	001	Oct 17, 2007
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CONS

SOLUTION;TOPICAL

DICLOFENAC SODIUM

AT		AMNEAL PHARMS	1.5%	A206116	001	Sep 02, 2016
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AT	!	APOTEX INC	1.5%	A202027	001	May 27, 2014
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AT		CADILA	1.5%	A206411	001	Apr 17, 2018
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AT		LUPIN LTD	1.5%	A204132	001	Aug 20, 2015
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AT		NOVEL LABS INC	1.5%	A205878	001	Dec 09, 2015
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AT		TARO	1.5%	A203818	001	Nov 26, 2014
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AT		TELLIGENT PHARMA INC	1.5%	A202769	001	Jul 08, 2015
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AT		TWI PHARMS	1.5%	A202393	001	Nov 24, 2014
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AT		WATSON LABS INC	1.5%	A202852	001	Nov 24, 2014
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PENNSAID

+! HORIZON

2%

				N204623	001	Jan 16, 2014
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SOLUTION/DROPS;OPHTHALMIC

DICLOFENAC SODIUM

AT		AKORN	0.1%	A077845	001	Apr 17, 2008
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AT		ALTAIRE PHARMS INC	0.1%	A203383	001	Nov 16, 2015
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AT		BAUSCH AND LOMB	0.1%	A078792	001	Dec 28, 2007
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AT		RISING	0.1%	A078553	001	Dec 28, 2007
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AT		SANDOZ INC	0.1%	A078031	001	Feb 06, 2008
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VOLTAREN

AT	+ !	NOVARTIS	0.1%	N020037	001	Mar 28, 1991
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TABLET, DELAYED RELEASE;ORAL

DICLOFENAC SODIUM

AB		ACTAVIS ELIZABETH	50MG	A074514	001	Mar 26, 1996
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AB			75MG	A074514	002	Mar 26, 1996
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AB		CARLSBAD	25MG	A075185	002	Nov 13, 1998
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AB			50MG	A075185	003	Nov 13, 1998
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AB			75MG	A075185	001	Nov 13, 1998
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AB		MYLAN PHARMS INC	50MG	A075281	002	Feb 12, 2002
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AB			75MG	A075281	003	Feb 12, 2002
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AB	!	UNIQUE PHARM LABS	25MG	A090066	001	Dec 01, 2010
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AB	!		50MG	A090066	002	Dec 01, 2010
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AB	!		75MG	A077863	003	Jun 08, 2007
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TABLET, EXTENDED RELEASE;ORAL

DICLOFENAC SODIUM

AB	!	DEXCEL LTD	100MG	A076201	001	Nov 06, 2002
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AB		VPNA	100MG	A075492	001	Feb 11, 2000
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PRESCRIPTION DRUG PRODUCT LIST

DICLOFENAC SODIUM; MISOPROSTOL

TABLET, DELAYED RELEASE;ORAL

ARTHROTEC

AB	+	GD SEARLE LLC	50MG;0.2MG	N020607 001	Dec 24, 1997
AB	+	!	75MG;0.2MG	N020607 002	Dec 24, 1997

DICLOFENAC SODIUM AND MISOPROSTOL

AB		ACTAVIS LABS FL INC	50MG;0.2MG	A201089 001	Jul 09, 2012
AB			75MG;0.2MG	A201089 002	Jul 09, 2012
AB		AMNEAL PHARMS	50MG;0.2MG	A203995 001	Nov 25, 2016
AB			75MG;0.2MG	A203995 002	Nov 25, 2016
AB		SANDOZ	50MG;0.2MG	A200158 001	May 09, 2013
AB			75MG;0.2MG	A200158 002	May 09, 2013
AB		YUNG SHIN PHARM	50MG;0.2MG	A205143 001	Feb 19, 2020
AB			75MG;0.2MG	A205143 002	Feb 19, 2020

DICLOXACILLIN SODIUM

CAPSULE;ORAL

DICLOXACILLIN SODIUM

AB		SANDOZ	EQ 250MG BASE	A061454 001	
AB		!	EQ 500MG BASE	A061454 003	
AB		TEVA	EQ 250MG BASE	A062286 001	Jun 03, 1982
AB			EQ 500MG BASE	A062286 002	Jun 03, 1982
		SANDOZ	EQ 125MG BASE	A061454 002	

DICYCLOMINE HYDROCHLORIDE

CAPSULE;ORAL

DICYCLOMINE HYDROCHLORIDE

AB		!	LANNETT	10MG	A084285 001
AB			MYLAN	10MG	A040319 001
AB			WATSON LABS	10MG	A085082 001
AB			WEST WARD	10MG	A040204 001

INJECTABLE;INJECTION

BENTYL

AP	+	!	ALLERGAN	10MG/ML	N008370 001	Oct 15, 1984
			BENTYL PRESERVATIVE FREE			
AP	+	!	ALLERGAN	10MG/ML	N008370 002	Oct 15, 1984

DICYCLOMINE HYDROCHLORIDE

AP		AKORN INC	10MG/ML	A207084 001	May 04, 2018
AP		AM REGENT	10MG/ML	A208353 001	Feb 17, 2017
AP		CUSTOPHARM INC	10MG/ML	A210788 001	Feb 11, 2019
AP		FOSUN PHARMA	10MG/ML	A210979 001	Jul 02, 2018
AP		FRESENIUS KABI USA	10MG/ML	A210257 001	Jan 25, 2019
AP		NEXUS PHARMS	10MG/ML	A206468 001	Feb 01, 2019
AP		SLATE	10MG/ML	A207076 001	Nov 02, 2018
AP		SUNGEN PHARMA	10MG/ML	A212058 001	Apr 26, 2019

DICYCLOMINE HYDROCHLORIDE (PRESERVATIVE FREE)

AP		WEST-WARD PHARMS	10MG/ML	A040465 001	Jun 30, 2003
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SYRUP;ORAL

DICYCLOMINE HYDROCHLORIDE

!	GENERIC	10MG/5ML	A040169 001	Mar 24, 2005
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TABLET;ORAL

DICYCLOMINE HYDROCHLORIDE

AB		HIKMA PHARMS	20MG	A040161 001	Oct 01, 1996	
AB		LANNETT	20MG	A040230 001	Feb 26, 1999	
AB		MYLAN	20MG	A040317 001	Sep 07, 1999	
AB		!	WATSON LABS	20MG	A085223 001	Jul 30, 1986

DIDANOSINE

CAPSULE, DELAYED REL PELLETS;ORAL

DIDANOSINE

AB		AUROBINDO PHARMA	125MG	A090094 001	Sep 24, 2008
AB			200MG	A090094 002	Sep 24, 2008
AB			250MG	A090094 003	Sep 24, 2008
AB			400MG	A090094 004	Sep 24, 2008

VIDEX EC

AB	+	BRISTOL MYERS	125MG	N021183 001	Oct 31, 2000
		SQUIBB			
AB	+		200MG	N021183 002	Oct 31, 2000
AB	+		250MG	N021183 003	Oct 31, 2000
AB	+	!	400MG	N021183 004	Oct 31, 2000

FOR SOLUTION;ORAL

VIDEX

+	!	BRISTOL-MYERS	10MG/ML	N020156 001	Oct 09, 1991
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PRESCRIPTION DRUG PRODUCT LIST

DIDANOSINE

FOR SOLUTION;ORAL
VIDEX
SQUIBB

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

AB ! LANNETT CO INC **75MG** **A091680 001** Oct 24, 2011

DIFLORASONE DIACETATE

CREAM;TOPICAL
DIFLORASONE DIACETATE

BX ! FOUGERA PHARMS 0.05% A076263 001 Dec 20, 2002
BX ! TARO 0.05% A075508 001 Apr 24, 2000

OINTMENT;TOPICAL

DIFLORASONE DIACETATE

AB AKORN **0.05%** **A206572 001** Jul 24, 2015
AB FOUGERA 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 TELIGENT PHARMA INC **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

DIFLUPREDNATE

EMULSION;OPHTHALMIC
DUREZOL

+! NOVARTIS 0.05% N022212 001 Jun 23, 2008

DIGOXIN

ELIXIR;ORAL

DIGOXIN

AA +! HIKMA **0.05MG/ML** **N021648 001** Aug 26, 2004
AA VISTAPHARM **0.05MG/ML** **A213000 001** Oct 04, 2019

INJECTABLE;INJECTION

DIGOXIN

AP SANDOZ INC **0.25MG/ML** **A040481 001** Aug 21, 2003
AP WEST-WARD PHARMS **0.25MG/ML** **A083391 001**

INT

LANOXIN

AP +! COVIS PHARMA BV **0.25MG/ML** **N009330 002**

LANOXIN PEDIATRIC

+! COVIS PHARMA BV 0.1MG/ML N009330 004

TABLET;ORAL

DIGOXIN

AB HIKMA INTL PHARMS **0.125MG** **A077002 002** Oct 30, 2007
AB **0.25MG** **A077002 001** Oct 30, 2007
AB IMPAX LABS **0.125MG** **A078556 001** Jul 20, 2009
AB **0.25MG** **A078556 002** Jul 20, 2009
AB MYLAN PHARMS INC **0.125MG** **A040282 001** Dec 23, 1999
AB **0.25MG** **A040282 002** Dec 23, 1999
AB STEVENS J **0.125MG** **A076268 001** Jul 26, 2002
AB **0.25MG** **A076268 002** Jul 26, 2002
AB SUN PHARM INDS INC **0.125MG** **A076363 001** Jan 31, 2003
AB **0.25MG** **A076363 002** Jan 31, 2003

LANOXIN

AB + CONCORDIA **0.125MG** **N020405 002** Sep 30, 1997
AB +! **0.25MG** **N020405 004** Sep 30, 1997
+ 0.0625MG N020405 001 Sep 30, 1997

PRESCRIPTION DRUG PRODUCT LIST

DIHYDROERGOTAMINE MESYLATE

INJECTABLE; INJECTION

D.H.E. 45

<u>AP</u>	<u>+!</u> BAUSCH	<u>1MG/ML</u>	<u>N005929</u>	<u>001</u>	
<u>DIHYDROERGOTAMINE MESYLATE</u>					
<u>AP</u>	APOLLO	<u>1MG/ML</u>	<u>A212046</u>	<u>001</u>	Jan 07, 2020
<u>AP</u>	HIKMA PHARMS	<u>1MG/ML</u>	<u>A206621</u>	<u>001</u>	Sep 15, 2017
<u>AP</u>	PADDOCK LLC	<u>1MG/ML</u>	<u>A040475</u>	<u>001</u>	Apr 28, 2003
<u>AP</u>	SAGENT PHARMS INC	<u>1MG/ML</u>	<u>A207264</u>	<u>001</u>	Jul 11, 2018
<u>AP</u>	WEST-WARD PHARMS INT	<u>1MG/ML</u>	<u>A040453</u>	<u>001</u>	Jun 09, 2003

SPRAY, METERED; NASAL

DIHYDROERGOTAMINE MESYLATE

<u>AB</u>	CUSTOPHARM INC	<u>0.5MG/SPRAY</u>	<u>A211393</u>	<u>001</u>	Feb 28, 2020
<u>MIGRANAL</u>					
<u>AB</u>	<u>+!</u> BAUSCH	<u>0.5MG/SPRAY</u>	<u>N020148</u>	<u>001</u>	Dec 08, 1997

DILTIAZEM HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

DILTIAZEM HYDROCHLORIDE

<u>AB2</u>	APOTEX	<u>180MG</u>	<u>A074943</u>	<u>002</u>	Dec 19, 2000
<u>AB2</u>	<u>!</u>	<u>240MG</u>	<u>A074943</u>	<u>001</u>	Aug 06, 1998

CARDIZEM CD

<u>AB3</u>	<u>+</u> BAUSCH	<u>120MG</u>	<u>N020062</u>	<u>001</u>	Aug 10, 1992
<u>AB3</u>	<u>+</u>	<u>180MG</u>	<u>N020062</u>	<u>002</u>	Dec 27, 1991
<u>AB3</u>	<u>+</u>	<u>240MG</u>	<u>N020062</u>	<u>003</u>	Dec 27, 1991
<u>AB3</u>	<u>+</u>	<u>300MG</u>	<u>N020062</u>	<u>004</u>	Dec 27, 1991
<u>AB3</u>	<u>+!</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>	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>	PAR PHARM	<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>	SUN PHARM	<u>120MG</u>	<u>A090492</u>	<u>001</u>	Oct 28, 2011
<u>AB3</u>		<u>120MG</u>	<u>A203023</u>	<u>001</u>	Jun 08, 2017
<u>AB3</u>		<u>180MG</u>	<u>A090492</u>	<u>002</u>	Oct 28, 2011
<u>AB3</u>		<u>180MG</u>	<u>A203023</u>	<u>002</u>	Jun 08, 2017
<u>AB3</u>		<u>240MG</u>	<u>A090492</u>	<u>003</u>	Oct 28, 2011
<u>AB3</u>		<u>240MG</u>	<u>A203023</u>	<u>003</u>	Jun 08, 2017
<u>AB3</u>		<u>300MG</u>	<u>A090492</u>	<u>004</u>	Oct 28, 2011
<u>AB3</u>		<u>300MG</u>	<u>A203023</u>	<u>004</u>	Jun 08, 2017
<u>AB3</u>		<u>360MG</u>	<u>A090492</u>	<u>005</u>	Oct 28, 2011
<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>	SANDOZ	<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

PRESCRIPTION DRUG PRODUCT LIST

DILTIAZEM HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE;ORAL

DILTIAZEM HYDROCHLORIDE

<u>AB4</u>		<u>360MG</u>	<u>A091022 005</u>	Sep 28, 2012
<u>AB4</u>		<u>420MG</u>	<u>A091022 006</u>	Sep 28, 2012
<u>AB4</u>	SUN PHARM	<u>120MG</u>	<u>A090421 001</u>	Nov 15, 2010
<u>AB4</u>		<u>180MG</u>	<u>A090421 002</u>	Nov 15, 2010
<u>AB4</u>		<u>240MG</u>	<u>A090421 003</u>	Nov 15, 2010
<u>AB4</u>		<u>300MG</u>	<u>A090421 004</u>	Nov 15, 2010
<u>AB4</u>		<u>360MG</u>	<u>A090421 005</u>	Nov 15, 2010
<u>AB4</u>	ZYDUS PHARMS	<u>120MG</u>	<u>A206641 001</u>	Aug 11, 2017
<u>AB4</u>		<u>180MG</u>	<u>A206641 002</u>	Aug 11, 2017
<u>AB4</u>		<u>240MG</u>	<u>A206641 003</u>	Aug 11, 2017
<u>AB4</u>		<u>300MG</u>	<u>A206641 004</u>	Aug 11, 2017
<u>AB4</u>		<u>360MG</u>	<u>A206641 005</u>	Aug 11, 2017
<u>AB4</u>		<u>420MG</u>	<u>A206641 006</u>	Aug 11, 2017

DILTIZAC

<u>AB4</u>	APOTEX INC	<u>120MG</u>	<u>A076395 001</u>	Feb 01, 2006
<u>AB4</u>		<u>180MG</u>	<u>A076395 002</u>	Feb 01, 2006
<u>AB4</u>		<u>240MG</u>	<u>A076395 003</u>	Feb 01, 2006
<u>AB4</u>		<u>300MG</u>	<u>A076395 004</u>	Feb 01, 2006
<u>AB4</u>		<u>360MG</u>	<u>A076395 005</u>	Feb 01, 2006

TAZTIA XT

<u>AB4</u>	ACTAVIS LABS FL INC	<u>120MG</u>	<u>A075401 001</u>	Apr 10, 2003
<u>AB4</u>		<u>180MG</u>	<u>A075401 002</u>	Apr 10, 2003
<u>AB4</u>		<u>240MG</u>	<u>A075401 003</u>	Apr 10, 2003
<u>AB4</u>		<u>300MG</u>	<u>A075401 004</u>	Apr 10, 2003
<u>AB4</u>		<u>360MG</u>	<u>A075401 005</u>	Apr 10, 2003

TIAZAC

<u>AB4</u>	+	BAUSCH	<u>120MG</u>	<u>N020401 001</u>	Sep 11, 1995
<u>AB4</u>	+		<u>180MG</u>	<u>N020401 002</u>	Sep 11, 1995
<u>AB4</u>	+		<u>240MG</u>	<u>N020401 003</u>	Sep 11, 1995
<u>AB4</u>	+		<u>300MG</u>	<u>N020401 004</u>	Sep 11, 1995
<u>AB4</u>	+		<u>360MG</u>	<u>N020401 005</u>	Sep 11, 1995
<u>AB4</u>	+	!	<u>420MG</u>	<u>N020401 006</u>	Oct 16, 1998

DILTIAZEM HYDROCHLORIDE

BC	!	MYLAN	120MG	A074910 003	May 02, 1997
		APOTEX	120MG	A074943 003	Dec 19, 2000
		MYLAN	60MG	A074910 001	May 02, 1997
			90MG	A074910 002	May 02, 1997

INJECTABLE; INJECTION

DILTIAZEM HYDROCHLORIDE

<u>AP</u>		AKORN INC	<u>5MG/ML</u>	<u>A075086 001</u>	Apr 09, 1998
<u>AP</u>	!	ATHENEX INC	<u>5MG/ML</u>	<u>A074617 001</u>	Feb 28, 1996
<u>AP</u>		HIKMA FARMACEUTICA	<u>5MG/ML</u>	<u>A202651 001</u>	Aug 09, 2012
<u>AP</u>		HOSPIRA	<u>5MG/ML</u>	<u>A074941 001</u>	Apr 15, 1998
<u>AP</u>		INTL MEDICATION	<u>5MG/ML</u>	<u>A075749 001</u>	Nov 21, 2001
<u>AP</u>		WEST-WARD PHARMS	<u>5MG/ML</u>	<u>A078538 001</u>	Dec 17, 2008
		INT			
	!	HOSPIRA	100MG/VIAL	A075853 001	Dec 17, 2002

TABLET; ORAL

CARDIZEM

<u>AB</u>	+	BAUSCH	<u>30MG</u>	<u>N018602 001</u>	Nov 05, 1982
<u>AB</u>	+		<u>60MG</u>	<u>N018602 002</u>	Nov 05, 1982
<u>AB</u>	+		<u>90MG</u>	<u>N018602 003</u>	Dec 08, 1986
<u>AB</u>	+	!	<u>120MG</u>	<u>N018602 004</u>	Dec 08, 1986

DILTIAZEM HYDROCHLORIDE

<u>AB</u>		EDENBRIDGE PHARMS	<u>30MG</u>	<u>A211596 001</u>	Nov 18, 2019
<u>AB</u>			<u>60MG</u>	<u>A211596 002</u>	Nov 18, 2019
<u>AB</u>			<u>90MG</u>	<u>A211596 003</u>	Nov 18, 2019
<u>AB</u>			<u>120MG</u>	<u>A211596 004</u>	Nov 18, 2019
<u>AB</u>		MYLAN	<u>30MG</u>	<u>A072838 004</u>	Nov 05, 1992
<u>AB</u>			<u>60MG</u>	<u>A072838 003</u>	Nov 05, 1992
<u>AB</u>			<u>90MG</u>	<u>A072838 002</u>	Nov 05, 1992
<u>AB</u>			<u>120MG</u>	<u>A072838 001</u>	Nov 05, 1992
<u>AB</u>		TEVA	<u>30MG</u>	<u>A074185 001</u>	May 31, 1995
<u>AB</u>			<u>60MG</u>	<u>A074185 002</u>	May 31, 1995
<u>AB</u>			<u>90MG</u>	<u>A074185 003</u>	May 31, 1995
<u>AB</u>			<u>120MG</u>	<u>A074185 004</u>	May 31, 1995

TABLET, EXTENDED RELEASE;ORAL

CARDIZEM LA

<u>AB</u>	+	BAUSCH	<u>120MG</u>	<u>N021392 001</u>	Feb 06, 2003
<u>AB</u>	+		<u>180MG</u>	<u>N021392 002</u>	Feb 06, 2003

PRESCRIPTION DRUG PRODUCT LIST

3-141 (of 453)

DILTIAZEM HYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL

CARDIZEM LA

AB	+	<u>240MG</u>	<u>N021392</u>	<u>003</u>	Feb 06, 2003
AB	+	<u>300MG</u>	<u>N021392</u>	<u>004</u>	Feb 06, 2003
AB	+	<u>360MG</u>	<u>N021392</u>	<u>005</u>	Feb 06, 2003
AB	+	<u>420MG</u>	<u>N021392</u>	<u>006</u>	Feb 06, 2003

DILTIAZEM HYDROCHLORIDE

AB	ACTAVIS LABS FL INC	<u>120MG</u>	<u>A077686</u>	<u>006</u>	Mar 15, 2010
AB		<u>180MG</u>	<u>A077686</u>	<u>005</u>	Mar 15, 2010
AB		<u>240MG</u>	<u>A077686</u>	<u>004</u>	Mar 15, 2010
AB		<u>300MG</u>	<u>A077686</u>	<u>003</u>	Mar 15, 2010
AB		<u>360MG</u>	<u>A077686</u>	<u>002</u>	Mar 15, 2010
AB		<u>420MG</u>	<u>A077686</u>	<u>001</u>	Mar 15, 2010

DIMENHYDRINATE

INJECTABLE; INJECTION

DIMENHYDRINATE

!	FRESENIUS KABI USA	50MG/ML	A040519	001	Jun 23, 2004
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DIMERCAPROL

INJECTABLE; INJECTION

BAL

+	AKORN	10%	N005939	001	
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DIMETHYL FUMARATE

CAPSULE, DELAYED RELEASE;ORAL

TECFIDERA

+	BIOGEN IDEC INC	120MG	N204063	001	Mar 27, 2013
+	!	240MG	N204063	002	Mar 27, 2013

DIMETHYL SULFOXIDE

SOLUTION; INTRAVESICAL

RIMSO-50

+	MYLAN INSTITUTIONAL	50%	N017788	001	
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DINOPROSTONE

GEL; ENDOCERVICAL

PREPIDIL

+	PHARMACIA AND UPJOHN	0.5MG/3GM	N019617	001	Dec 09, 1992
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INSERT, EXTENDED RELEASE; VAGINAL

CERVIDIL

+	FERRING PHARMS INC	10MG	N020411	001	Mar 30, 1995
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SUPPOSITORY; VAGINAL

PROSTIN E2

+	PHARMACIA AND UPJOHN	20MG	N017810	001	
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DIPHENHYDRAMINE HYDROCHLORIDE

ELIXIR; ORAL

DIPHENHYDRAMINE HYDROCHLORIDE

!	PHARM ASSOC	12.5MG/5ML	A087513	001	Feb 10, 1982
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INJECTABLE; INJECTION

DIPHENHYDRAMINE HYDROCHLORIDE

AP	APP PHARMS	<u>50MG/ML</u>	<u>A040466</u>	<u>001</u>	May 28, 2002
AP	HOSPIRA	<u>50MG/ML</u>	<u>A040140</u>	<u>001</u>	Nov 20, 1998
AP	MICRO LABS	<u>50MG/ML</u>	<u>A205723</u>	<u>001</u>	Aug 22, 2018
AP	MYLAN INSTITUTIONAL	<u>50MG/ML</u>	<u>A040498</u>	<u>001</u>	Jul 12, 2005
AP	!	<u>50MG/ML</u>	<u>A080817</u>	<u>002</u>	

DIPHENHYDRAMINE HYDROCHLORIDE PRESERVATIVE FREE

AP	FRESENIUS KABI USA	<u>50MG/ML</u>	<u>A091526</u>	<u>001</u>	Mar 26, 2013
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DIPYRIDAMOLE

INJECTABLE; INJECTION

DIPYRIDAMOLE

AP	!	ATHENEX INC	<u>5MG/ML</u>	<u>A074939</u>	<u>001</u>	Apr 13, 1998
AP		FRESENIUS KABI USA	<u>5MG/ML</u>	<u>A074956</u>	<u>001</u>	Sep 30, 1998
AP		WEST-WARD PHARMS	<u>5MG/ML</u>	<u>A074521</u>	<u>001</u>	Oct 18, 1996

TABLET; ORAL

DIPYRIDAMOLE

AB	BARR	<u>25MG</u>	<u>A087184</u>	<u>001</u>	Oct 03, 1990
AB		<u>50MG</u>	<u>A087716</u>	<u>001</u>	Oct 03, 1990
AB		<u>75MG</u>	<u>A087717</u>	<u>001</u>	Oct 03, 1990
AB	IMPAX LABS	<u>25MG</u>	<u>A040782</u>	<u>001</u>	Jul 18, 2007

PRESCRIPTION DRUG PRODUCT LIST

DIPYRIDAMOLE

TABLET;ORAL

DIPYRIDAMOLE

<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>	MURTY PHARMS	<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 INC	<u>25MG</u>	<u>A040874</u>	<u>001</u>	Jan 28, 2008
<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
<u>PERSANTINE</u>					
<u>AB</u>	+ BOEHRINGER INGELHEIM	<u>25MG</u>	<u>N012836</u>	<u>003</u>	Dec 22, 1986
<u>AB</u>	+	<u>50MG</u>	<u>N012836</u>	<u>004</u>	Feb 06, 1987
<u>AB</u>	+	<u>75MG</u>	<u>N012836</u>	<u>005</u>	Feb 06, 1987

DIROXIMEL FUMARATE

CAPSULE, DELAYED RELEASE;ORAL

VUMERITY

+! BIOGEN 231MG N211855 001 Oct 29, 2019

DISOPYRAMIDE PHOSPHATE

CAPSULE;ORAL

DISOPYRAMIDE PHOSPHATE

<u>AB</u>	MAYNE PHARMA	<u>EQ 100MG BASE</u>	<u>A070173</u>	<u>001</u>	May 31, 1985
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A070173</u>	<u>002</u>	May 31, 1985
<u>AB</u>	TEVA	<u>EQ 100MG BASE</u>	<u>A070101</u>	<u>001</u>	Feb 22, 1985
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A070102</u>	<u>001</u>	Feb 22, 1985

NORPACE

<u>AB</u>	+ GD SEARLE LLC	<u>EQ 100MG BASE</u>	<u>N017447</u>	<u>001</u>	
<u>AB</u>	+	<u>EQ 150MG BASE</u>	<u>N017447</u>	<u>002</u>	
CAPSULE, EXTENDED RELEASE;ORAL					
NORPACE CR					
	+ GD SEARLE LLC	<u>EQ 100MG BASE</u>	N018655	001	Jul 20, 1982
	+	<u>EQ 150MG BASE</u>	N018655	002	Jul 20, 1982

DISULFIRAM

TABLET;ORAL

ANTABUSE

<u>AB</u>	ODYSSEY PHARMS	<u>250MG</u>	<u>A088482</u>	<u>001</u>	Dec 08, 1983
<u>AB</u>	!	<u>500MG</u>	<u>A088483</u>	<u>001</u>	Dec 08, 1983

DISULFIRAM

<u>AB</u>	ALVOGEN	<u>250MG</u>	<u>A091681</u>	<u>001</u>	Aug 08, 2013
<u>AB</u>	CHARTWELL MOLECULES	<u>250MG</u>	<u>A091563</u>	<u>001</u>	Dec 31, 2012
<u>AB</u>		<u>500MG</u>	<u>A091563</u>	<u>002</u>	Dec 31, 2012
<u>AB</u>	SIGMAPHARM LABS LLC	<u>250MG</u>	<u>A091619</u>	<u>001</u>	Mar 28, 2011
<u>AB</u>		<u>500MG</u>	<u>A091619</u>	<u>002</u>	Mar 28, 2011

DIVALPROEX SODIUM

CAPSULE, DELAYED REL PELLETS;ORAL

DEPAKOTE

<u>AB</u>	+! ABBVIE	<u>EQ 125MG VALPROIC ACID</u>	<u>N019680</u>	<u>001</u>	Sep 12, 1989
<u>DIVALPROEX SODIUM</u>					
<u>AB</u>	AJANTA PHARMA LTD	<u>EQ 125MG VALPROIC ACID</u>	<u>A213181</u>	<u>001</u>	Mar 02, 2020
<u>AB</u>	DR REDDYS LABS LTD	<u>EQ 125MG VALPROIC ACID</u>	<u>A078979</u>	<u>001</u>	Jan 23, 2009
<u>AB</u>	ZYDUS PHARMS USA INC	<u>EQ 125MG VALPROIC ACID</u>	<u>A078919</u>	<u>001</u>	Jan 27, 2009

TABLET, DELAYED RELEASE;ORAL

DEPAKOTE

<u>AB</u>	+ ABBVIE	<u>EQ 125MG VALPROIC ACID</u>	<u>N018723</u>	<u>003</u>	Oct 26, 1984
<u>AB</u>	+	<u>EQ 250MG VALPROIC ACID</u>	<u>N018723</u>	<u>001</u>	Mar 10, 1983
<u>AB</u>	+	<u>EQ 500MG VALPROIC ACID</u>	<u>N018723</u>	<u>002</u>	Mar 10, 1983

DIVALPROEX SODIUM

<u>AB</u>	ACTAVIS LABS FL INC	<u>EQ 500MG VALPROIC ACID</u>	<u>A079080</u>	<u>001</u>	Feb 25, 2011
<u>AB</u>	ANCHEN PHARMS	<u>EQ 500MG VALPROIC ACID</u>	<u>A078411</u>	<u>001</u>	Nov 03, 2008
<u>AB</u>	APOTEX	<u>EQ 125MG VALPROIC ACID</u>	<u>A077615</u>	<u>003</u>	Jul 29, 2008
<u>AB</u>		<u>EQ 250MG VALPROIC ACID</u>	<u>A077615</u>	<u>002</u>	Jul 29, 2008
<u>AB</u>		<u>EQ 500MG VALPROIC ACID</u>	<u>A077615</u>	<u>001</u>	Jul 29, 2008
<u>AB</u>	AUROBINDO PHARMA LTD	<u>EQ 125MG VALPROIC ACID</u>	<u>A090554</u>	<u>001</u>	Apr 21, 2011
<u>AB</u>		<u>EQ 250MG VALPROIC ACID</u>	<u>A090554</u>	<u>002</u>	Apr 21, 2011
<u>AB</u>		<u>EQ 500MG VALPROIC ACID</u>	<u>A090554</u>	<u>003</u>	Apr 21, 2011
<u>AB</u>	CELLTRION	<u>EQ 125MG VALPROIC ACID</u>	<u>A077296</u>	<u>001</u>	Jul 31, 2008

PRESCRIPTION DRUG PRODUCT LIST

DIVALPROEX SODIUM

TABLET, DELAYED RELEASE;ORAL

DIVALPROEX SODIUM

<u>AB</u>		<u>EQ 250MG VALPROIC ACID</u>	<u>A077296 002</u>	Jul 31, 2008
<u>AB</u>		<u>EQ 500MG VALPROIC ACID</u>	<u>A077296 003</u>	Jul 31, 2008
<u>AB</u>	DR REDDYS LABS LTD	<u>EQ 125MG VALPROIC ACID</u>	<u>A078755 001</u>	Jul 29, 2008
<u>AB</u>		<u>EQ 250MG VALPROIC ACID</u>	<u>A078755 002</u>	Jul 29, 2008
<u>AB</u>		<u>EQ 500MG VALPROIC ACID</u>	<u>A078755 003</u>	Jul 29, 2008
<u>AB</u>	INVATECH	<u>EQ 125MG VALPROIC ACID</u>	<u>A078290 003</u>	Jul 29, 2008
<u>AB</u>		<u>EQ 250MG VALPROIC ACID</u>	<u>A078290 002</u>	Jul 29, 2008
<u>AB</u>		<u>EQ 500MG VALPROIC ACID</u>	<u>A078290 001</u>	Jul 29, 2008
<u>AB</u>	LUPIN	<u>EQ 125MG VALPROIC ACID</u>	<u>A078790 001</u>	Jul 29, 2008
<u>AB</u>		<u>EQ 250MG VALPROIC ACID</u>	<u>A078790 002</u>	Jul 29, 2008
<u>AB</u>		<u>EQ 500MG VALPROIC ACID</u>	<u>A078790 003</u>	Jul 29, 2008
<u>AB</u>	ORCHID HLTHCARE	<u>EQ 125MG VALPROIC ACID</u>	<u>A078853 001</u>	Nov 25, 2008
<u>AB</u>		<u>EQ 250MG VALPROIC ACID</u>	<u>A078853 002</u>	Nov 25, 2008
<u>AB</u>		<u>EQ 500MG VALPROIC ACID</u>	<u>A078853 003</u>	Nov 25, 2008
<u>AB</u>	PRINSTON INC	<u>EQ 125MG VALPROIC ACID</u>	<u>A090210 001</u>	Nov 30, 2009
<u>AB</u>		<u>EQ 250MG VALPROIC ACID</u>	<u>A090210 002</u>	Nov 30, 2009
<u>AB</u>		<u>EQ 500MG VALPROIC ACID</u>	<u>A090210 003</u>	Nov 30, 2009
<u>AB</u>	SUN PHARM INDS	<u>EQ 125MG VALPROIC ACID</u>	<u>A078597 001</u>	Jul 29, 2008
<u>AB</u>		<u>EQ 250MG VALPROIC ACID</u>	<u>A078597 002</u>	Jul 29, 2008
<u>AB</u>		<u>EQ 500MG VALPROIC ACID</u>	<u>A078597 003</u>	Jul 29, 2008
<u>AB</u>	UNICHEM LABS LTD	<u>EQ 125MG VALPROIC ACID</u>	<u>A079163 001</u>	Apr 05, 2011
<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>	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>	IMPAX LABS	<u>EQ 250MG VALPROIC ACID</u>	<u>A078791 001</u>	May 06, 2009
<u>AB</u>		<u>EQ 500MG VALPROIC ACID</u>	<u>A078791 002</u>	Aug 04, 2009
<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>	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>	HOSPIRA	<u>EQ 12.5MG BASE/ML</u>	<u>A074086 001</u>	Nov 29, 1993
<u>AP</u>	!	<u>EQ 12.5MG BASE/ML</u>	<u>A074292 001</u>	Feb 16, 1995
<u>AP</u>	WEST-WARD PHARMS INT	<u>EQ 12.5MG BASE/ML</u>	<u>A074277 001</u>	Oct 31, 1994

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

PRESCRIPTION DRUG PRODUCT LIST

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 LLC	<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>		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>		DFB ONCOLOGY LTD	<u>20MG/ML (20MG/ML)</u>	<u>A206177 001</u>	Jan 20, 2017
<u>AP</u>			<u>80MG/4ML (20MG/ML)</u>	<u>A206177 002</u>	Jan 20, 2017
<u>AP</u>		DR REDDYS LABS LTD	<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>	+	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>		INGENUS PHARMS LLC	<u>20MG/2ML (10MG/ML)</u>	<u>A207563 001</u>	Aug 31, 2017
<u>AP</u>			<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>		JIANGSU HENGRUI MED	<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>		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>		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 MEDICARE LTD	<u>20MG/ML (20MG/ML)</u>	<u>A210327 001</u>	May 16, 2019
<u>AP</u>			<u>80MG/4ML (20MG/ML)</u>	<u>A210327 002</u>	May 16, 2019
<u>AP</u>			<u>160MG/8ML (20MG/ML)</u>	<u>A210327 003</u>	May 16, 2019
<u>AP</u>		TEIKOKU PHARMA	<u>20MG/ML (20MG/ML)</u>	<u>N205934 001</u>	Dec 22, 2015
<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>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 LLC	140MG/7ML (20MG/ML)	N203551 003	Apr 12, 2013
		DFB ONCOLOGY LTD	200MG/10ML (20MG/ML)	A206177 003	Jan 20, 2017
	+	HOSPIRA INC	20MG/ML (20MG/ML)	N022234 004	Jun 23, 2016
	+		80MG/4ML (20MG/ML)	N022234 005	Jun 23, 2016
	+		160MG/8ML (20MG/ML)	N022234 007	Jan 24, 2017
	!	JIANGSU HENGRUI MED	40MG/ML	A203170 001	Feb 15, 2017

SOLUTION; INTRAVENOUS

DOCETAXEL

<u>AP</u>		SUN PHARMA GLOBAL	<u>20MG/ML (20MG/ML)</u>	<u>N022534 003</u>	Jan 08, 2019
<u>AP</u>			<u>80MG/4ML (20MG/ML)</u>	<u>N022534 004</u>	Jan 08, 2019
<u>AP</u>			<u>160MG/8ML (20MG/ML)</u>	<u>N022534 005</u>	Jan 08, 2019

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>		BIONPHARMA INC	<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>		MAYNE PHARMA INC	<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>		MSN	<u>0.125MG</u>	<u>A213220 001</u>	Jan 29, 2020
<u>AB</u>			<u>0.25MG</u>	<u>A213220 002</u>	Jan 29, 2020
<u>AB</u>			<u>0.5MG</u>	<u>A213220 003</u>	Jan 29, 2020
<u>AB</u>		PAR PHARM INC	<u>0.125MG</u>	<u>A208519 001</u>	Oct 09, 2018

PRESCRIPTION DRUG PRODUCT LIST

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DOFETILIDE

CAPSULE;ORAL

DOFETILIDE

<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>	PRINSTON INC	<u>0.125MG</u>	<u>A211223</u>	<u>001</u>	Dec 17, 2019
<u>AB</u>		<u>0.25MG</u>	<u>A211223</u>	<u>002</u>	Dec 17, 2019
<u>AB</u>		<u>0.5MG</u>	<u>A211223</u>	<u>003</u>	Dec 17, 2019
<u>AB</u>	RICONPHARMA LLC	<u>0.125MG</u>	<u>A212410</u>	<u>001</u>	Dec 27, 2019
<u>AB</u>		<u>0.25MG</u>	<u>A212410</u>	<u>002</u>	Dec 27, 2019
<u>AB</u>		<u>0.5MG</u>	<u>A212410</u>	<u>003</u>	Dec 27, 2019
<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>	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 10MG BASE	N204790	002	Jun 09, 2016
+		EQ 25MG BASE	N204790	003	Jun 09, 2016
+	!	EQ 50MG BASE	N204790	001	Aug 12, 2013

DOLUTEGRAVIR SODIUM; LAMIVUDINE

TABLET;ORAL

DOVATO

+	VIIV HLTHCARE	EQ 50MG BASE;300MG	N211994	001	Apr 08, 2019
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DOLUTEGRAVIR SODIUM; RILPIVIRINE HYDROCHLORIDE

TABLET;ORAL

JULUCA

+	VIIV HLTHCARE	EQ 50MG BASE;EQ 25MG BASE	N210192	001	Nov 21, 2017
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DONEPEZIL HYDROCHLORIDE

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>		ACI HEALTHCARE LTD	<u>5MG</u>	<u>A078662</u>	<u>001</u>	May 31, 2011
<u>AB</u>			<u>10MG</u>	<u>A078662</u>	<u>002</u>	May 31, 2011
<u>AB</u>		ACTAVIS ELIZABETH	<u>23MG</u>	<u>A202415</u>	<u>001</u>	Dec 17, 2015
<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	<u>5MG</u>	<u>A090100</u>	<u>001</u>	Oct 24, 2012
<u>AB</u>			<u>10MG</u>	<u>A090100</u>	<u>002</u>	Oct 24, 2012
<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>		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>		CSPC OUYI	<u>5MG</u>	<u>A202114</u>	<u>001</u>	Jul 05, 2013
<u>AB</u>			<u>10MG</u>	<u>A202114</u>	<u>002</u>	Jul 05, 2013
<u>AB</u>		DEXCEL PHARMA	<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>		HERITAGE PHARMA	<u>5MG</u>	<u>A077344</u>	<u>001</u>	May 31, 2011
<u>AB</u>			<u>10MG</u>	<u>A077344</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>		HISUN PHARM	<u>23MG</u>	<u>A202410</u>	<u>001</u>	Mar 24, 2017
		HANGZHOU				
<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>			<u>23MG</u>	<u>A203419</u>	<u>001</u>	Apr 12, 2016
<u>AB</u>		JUBILANT GENERICS	<u>5MG</u>	<u>A090768</u>	<u>001</u>	May 31, 2011

PRESCRIPTION DRUG PRODUCT LIST

DONEPEZIL HYDROCHLORIDE

TABLET; ORAL

DONEPEZIL HYDROCHLORIDE

<u>AB</u>		<u>10MG</u>	<u>A090768 002</u>	May 31, 2011
<u>AB</u>	LUPIN LTD	<u>23MG</u>	<u>A202782 001</u>	Oct 30, 2015
<u>AB</u>	MACLEODS PHARMS LTD	<u>5MG</u>	<u>A201146 001</u>	Aug 17, 2012
<u>AB</u>		<u>10MG</u>	<u>A201146 002</u>	Aug 17, 2012
<u>AB</u>		<u>23MG</u>	<u>A202631 001</u>	Jan 22, 2014
<u>AB</u>	MYLAN	<u>23MG</u>	<u>A202656 001</u>	Oct 22, 2015
<u>AB</u>	MYLAN PHARMS INC	<u>5MG</u>	<u>A090521 001</u>	May 31, 2011
<u>AB</u>		<u>10MG</u>	<u>A090521 002</u>	May 31, 2011
<u>AB</u>	PLIVA HRVATSKA DOO	<u>5MG</u>	<u>A090425 001</u>	May 31, 2011
<u>AB</u>		<u>10MG</u>	<u>A090425 002</u>	May 31, 2011
<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>	SANDOZ	<u>5MG</u>	<u>A090290 001</u>	May 31, 2011
<u>AB</u>		<u>10MG</u>	<u>A090290 002</u>	May 31, 2011
<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>A090551 001</u>	May 31, 2011
<u>AB</u>		<u>10MG</u>	<u>A090551 002</u>	May 31, 2011
<u>AB</u>	SUN PHARM INDS	<u>5MG</u>	<u>A090493 001</u>	May 31, 2011
<u>AB</u>		<u>10MG</u>	<u>A090493 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>	UNICHEM LABS LTD	<u>5MG</u>	<u>A203656 001</u>	Jun 23, 2016
<u>AB</u>		<u>10MG</u>	<u>A203656 002</u>	Jun 23, 2016
<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
<u>AB</u>	SANDOZ	<u>5MG</u>	<u>A091198 001</u>	May 10, 2011
<u>AB</u>	!	<u>10MG</u>	<u>A091198 002</u>	May 10, 2011
<u>AB</u>	UNICHEM LABS LTD	<u>5MG</u>	<u>A204831 001</u>	Nov 10, 2016
<u>AB</u>		<u>10MG</u>	<u>A204831 002</u>	Nov 10, 2016

DONEPEZIL HYDROCHLORIDE; MEMANTINE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

NAMZARIC

+	ALLERGAN	10MG; 7MG	N206439 003	Jul 18, 2016
+		10MG; 14MG	N206439 001	Dec 23, 2014
+		10MG; 21MG	N206439 004	Jul 18, 2016
+	!	10MG; 28MG	N206439 002	Dec 23, 2014

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>	+!	<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 AND DEXTROSE 5%

<u>AP</u>	+!	B BRAUN	<u>80MG/100ML</u>	<u>N019099 002</u>	Oct 15, 1986
<u>AP</u>	+!		<u>320MG/100ML</u>	<u>N019099 004</u>	Oct 15, 1986

DOPAMINE HYDROCHLORIDE AND DEXTROSE 5% IN PLASTIC CONTAINER

<u>AP</u>	+!	B BRAUN	<u>160MG/100ML</u>	<u>N019099 003</u>	Oct 15, 1986
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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

DOPAMINE HYDROCHLORIDE AND DEXTROSE 5% IN PLASTIC CONTAINER

+	B BRAUN	40MG/100ML	N019099 001	Oct 15, 1986
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PRESCRIPTION DRUG PRODUCT LIST

DOPAMINE HYDROCHLORIDE

INJECTABLE; INJECTION

DOPAMINE HYDROCHLORIDE IN DEXTROSE 5% IN PLASTIC CONTAINER

+! 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 PHARMS LTD	<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>	HI TECH PHARMA	<u>EQ 2% BASE</u>	<u>A077846 001</u>	Oct 28, 2008
<u>AT</u>	MICRO LABS	<u>EQ 2% BASE</u>	<u>A204778 001</u>	Nov 08, 2019
<u>AT</u>	SANDOZ INC	<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

TRUSOPT

<u>AT</u>	+! MERCK	<u>EQ 2% BASE</u>	<u>N020408 001</u>	Dec 09, 1994
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DORZOLAMIDE HYDROCHLORIDE; TIMOLOL MALEATE

SOLUTION/DROPS; OPHTHALMIC

COSOPT

<u>AT</u>	+! OAK PHARMS INC	<u>EQ 2% BASE; EQ 0.5% BASE</u>	<u>N020869 001</u>	Apr 07, 1998
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COSOPT PF

<u>AT</u>	+! OAK PHARMS INC	<u>EQ 2% BASE; EQ 0.5% BASE</u>	<u>N202667 001</u>	Feb 01, 2012
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DORZOLAMIDE HYDROCHLORIDE AND TIMOLOL MALEATE

<u>AT</u>	AKORN INC	<u>EQ 2% BASE; EQ 0.5% BASE</u>	<u>A203058 001</u>	Sep 22, 2014
<u>AT</u>	AUROBINDO PHARMA LTD	<u>EQ 2% BASE; EQ 0.5% BASE</u>	<u>A207630 001</u>	Jul 24, 2018
<u>AT</u>	BAUSCH AND LOMB	<u>EQ 2% BASE; EQ 0.5% BASE</u>	<u>A090037 001</u>	Jul 14, 2009
<u>AT</u>	HI TECH PHARMA	<u>EQ 2% BASE; EQ 0.5% BASE</u>	<u>A077847 001</u>	Oct 28, 2008
<u>AT</u>	SANDOZ	<u>EQ 2% BASE; EQ 0.5% BASE</u>	<u>A078749 001</u>	Nov 06, 2008
<u>AT</u>	SANDOZ INC	<u>EQ 2% BASE; EQ 0.5% BASE</u>	<u>A090604 001</u>	Nov 18, 2009
<u>AT</u>	SOMERSET	<u>EQ 2% BASE; EQ 0.5% BASE</u>	<u>A207523 001</u>	Jun 25, 2019
<u>AT</u>	TEVA PHARMS	<u>EQ 2% BASE; EQ 0.5% BASE</u>	<u>A078704 001</u>	Sep 28, 2009

DOXAPRAM HYDROCHLORIDE

INJECTABLE; INJECTION

DOPRAM

<u>AP</u>	+! HIKMA	<u>20MG/ML</u>	<u>N014879 001</u>	
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DOXAPRAM HYDROCHLORIDE

<u>AP</u>	ATHENEX INC	<u>20MG/ML</u>	<u>A076266 001</u>	Jan 10, 2003
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DOXAZOSIN MESYLATE

TABLET; ORAL

CARDURA

<u>AB</u>	+! PFIZER	<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>	DAVA PHARMS INC	<u>EQ 1MG BASE</u>	<u>A076161 001</u>	Jun 10, 2004
<u>AB</u>		<u>EQ 2MG BASE</u>	<u>A076161 002</u>	Jun 10, 2004
<u>AB</u>		<u>EQ 4MG BASE</u>	<u>A076161 003</u>	Jun 10, 2004
<u>AB</u>		<u>EQ 8MG BASE</u>	<u>A076161 004</u>	Jun 10, 2004
<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

PRESCRIPTION DRUG PRODUCT LIST

DOXAZOSIN MESYLATE

TABLET;ORAL

DOXAZOSIN MESYLATE

<u>AB</u>	MYLAN	<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>	PLIVA	<u>EQ 1MG BASE</u>	<u>A075750 001</u>	Jun 08, 2001
<u>AB</u>		<u>EQ 2MG BASE</u>	<u>A075750 002</u>	Jun 08, 2001
<u>AB</u>		<u>EQ 4MG BASE</u>	<u>A075750 003</u>	Jun 08, 2001
<u>AB</u>		<u>EQ 8MG BASE</u>	<u>A075750 004</u>	Jun 08, 2001
<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>	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

+	PFIZER	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>	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>	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>	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>	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>	PAR PHARM	<u>EQ 10MG BASE</u>	<u>A071422 002</u>	Nov 09, 1987
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>A071422 003</u>	Nov 09, 1987
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A071422 004</u>	Nov 09, 1987
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A071422 005</u>	Nov 09, 1987
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A071422 001</u>	Nov 09, 1987
<u>AB</u>	!	EQ 150MG BASE	A071422 006	Nov 09, 1987

CONCENTRATE;ORAL

DOXEPIN HYDROCHLORIDE

<u>AA</u>	LANNETT CO INC	<u>EQ 10MG BASE/ML</u>	<u>A074721 001</u>	Dec 29, 1998
<u>AA</u>	!	<u>EQ 10MG BASE/ML</u>	<u>A071609 001</u>	Nov 09, 1987
<u>AA</u>	WOCKHARDT BIO AG	<u>EQ 10MG BASE/ML</u>	<u>A071918 001</u>	Jul 20, 1988

CREAM;TOPICAL

ZONALON

+	!	MYLAN	5%	N020126 001	Apr 01, 1994
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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

SILENOR

<u>AB</u>	+	CURRAX	<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

PRESCRIPTION DRUG PRODUCT LIST

DOXERCALCIFEROL

CAPSULE; ORAL

DOXERCALCIFEROL

<u>AB</u>	HIKMA	<u>0.5MCG</u>	<u>A091433 001</u>	Sep 23, 2011
<u>AB</u>		<u>1MCG</u>	<u>A091433 002</u>	Jan 14, 2014
<u>AB</u>		<u>2.5MCG</u>	<u>A091433 003</u>	Jan 14, 2014
<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

HECTOROL

<u>AB</u>	+ SANOFI	<u>0.5MCG</u>	<u>N020862 002</u>	Apr 23, 2004
<u>AB</u>	+	<u>1MCG</u>	<u>N020862 003</u>	Jul 13, 2009
<u>AB</u>	+	<u>2.5MCG</u>	<u>N020862 001</u>	Jun 09, 1999

INJECTABLE; INJECTION

DOXERCALCIFEROL

<u>AP</u>	AKORN INC	<u>2MCG/ML (2MCG/ML)</u>	<u>A203929 002</u>	Mar 28, 2016
<u>AP</u>		<u>4MCG/2ML (2MCG/ML)</u>	<u>A203929 001</u>	May 07, 2015
<u>AP</u>	AMNEAL	<u>2MCG/ML (2MCG/ML)</u>	<u>A208974 001</u>	May 24, 2017
<u>AP</u>		<u>4MCG/2ML (2MCG/ML)</u>	<u>A208974 002</u>	May 24, 2017
<u>AP</u>		<u>4MCG/2ML (2MCG/ML)</u>	<u>A208975 001</u>	May 24, 2017
<u>AP</u>	GLAND PHARMA LTD	<u>4MCG/2ML (2MCG/ML)</u>	<u>A210452 001</u>	Sep 26, 2019
<u>AP</u>	HIKMA PHARMS	<u>4MCG/2ML (2MCG/ML)</u>	<u>A091101 001</u>	Aug 30, 2013
<u>AP</u>	+ HOSPIRA INC	<u>4MCG/2ML (2MCG/ML)</u>	<u>N208614 001</u>	Jul 24, 2018
<u>AP</u>	LUPIN LTD	<u>4MCG/2ML (2MCG/ML)</u>	<u>A210801 001</u>	Nov 01, 2018
<u>AP</u>	MEITHEAL	<u>4MCG/2ML (2MCG/ML)</u>	<u>A211670 001</u>	Feb 07, 2020
<u>AP</u>	SANDOZ INC	<u>4MCG/2ML (2MCG/ML)</u>	<u>A091333 001</u>	May 05, 2014
<u>AP</u>		<u>4MCG/2ML (2MCG/ML)</u>	<u>A200926 001</u>	Feb 04, 2014
<u>AP</u>	SUN PHARM	<u>2MCG/ML (2MCG/ML)</u>	<u>A203875 001</u>	Nov 14, 2019
<u>AP</u>		<u>4MCG/2ML (2MCG/ML)</u>	<u>A203875 002</u>	Nov 14, 2019

HECTOROL

<u>AP</u>	+ SANOFI	<u>2MCG/ML (2MCG/ML)</u>	<u>N021027 002</u>	Apr 06, 2000
<u>AP</u>	+	<u>4MCG/2ML (2MCG/ML)</u>	<u>N021027 001</u>	Apr 06, 2000

DOXORUBICIN HYDROCHLORIDE

INJECTABLE; INJECTION

DOXORUBICIN HYDROCHLORIDE

<u>AP</u>	ACTAVIS INC	<u>2MG/ML</u>	<u>A203622 001</u>	Jun 27, 2014
<u>AP</u>		<u>200MG/100ML</u>	<u>A203622 002</u>	Jun 27, 2014
<u>AP</u>	AMNEAL	<u>20MG/VIAL</u>	<u>A208888 001</u>	Feb 17, 2017
<u>AP</u>		<u>50MG/VIAL</u>	<u>A208888 002</u>	Feb 17, 2017
<u>AP</u>	FRESENIUS KABI USA	<u>2MG/ML</u>	<u>A063277 001</u>	Oct 26, 1995
<u>AP</u>	GLAND PHARMA LTD	<u>2MG/ML</u>	<u>A209825 001</u>	Aug 11, 2017
<u>AP</u>	MYLAN LABS LTD	<u>50MG/VIAL</u>	<u>A200170 002</u>	Oct 28, 2011
<u>AP</u>	PHARMACHEMIE BV	<u>2MG/ML</u>	<u>A063336 001</u>	Feb 28, 1995
<u>AP</u>		<u>10MG/VIAL</u>	<u>A063097 001</u>	May 21, 1990
<u>AP</u>		<u>20MG/VIAL</u>	<u>A063097 002</u>	May 21, 1990
<u>AP</u>		<u>50MG/VIAL</u>	<u>A063097 003</u>	May 21, 1990
<u>AP</u>		<u>200MG/100ML</u>	<u>A063336 004</u>	Feb 28, 1995
<u>AP</u>	+! PHARMACIA AND UPJOHN	<u>2MG/ML</u>	<u>N050629 001</u>	Dec 23, 1987
<u>AP</u>	+!	<u>200MG/100ML</u>	<u>N050629 002</u>	May 03, 1988
<u>AP</u>	SAGENT PHARMS	<u>2MG/ML</u>	<u>A091495 001</u>	Mar 18, 2013
<u>AP</u>	SUN PHARM INDS	<u>2MG/ML</u>	<u>A091418 001</u>	Feb 15, 2012
<u>AP</u>	TEVA PHARMS USA	<u>2MG/ML</u>	<u>A064140 001</u>	Jul 28, 1995
<u>AP</u>		<u>200MG/100ML</u>	<u>A064140 002</u>	Jul 28, 1995
<u>AP</u>	WEST-WARD PHARMS INT	<u>2MG/ML</u>	<u>A062975 001</u>	Mar 17, 1989
<u>AP</u>	!	<u>10MG/VIAL</u>	<u>A062921 001</u>	Mar 17, 1989
<u>AP</u>	!	<u>20MG/VIAL</u>	<u>A062921 002</u>	Mar 17, 1989
<u>AP</u>	!	<u>50MG/VIAL</u>	<u>A062921 003</u>	Mar 17, 1989
<u>AP</u>		<u>200MG/100ML</u>	<u>A064097 001</u>	Sep 13, 1994
<u>AP</u>	+ PHARMACIA AND UPJOHN	150MG/75ML	N050629 003	Mar 28, 2011

INJECTABLE, LIPOSOMAL; INJECTION

DOXIL (LIPOSOMAL)

<u>AB</u>	+ BAXTER HLTHCARE CORP	<u>20MG/10ML (2MG/ML)</u>	<u>N050718 001</u>	Nov 17, 1995
<u>AB</u>	+	<u>50MG/25ML (2MG/ML)</u>	<u>N050718 002</u>	Jun 13, 2000

DOXORUBICIN HYDROCHLORIDE (LIPOSOMAL)

<u>AB</u>	DR REDDYS LABS LTD	<u>20MG/10ML (2MG/ML)</u>	<u>A208657 001</u>	May 15, 2017
<u>AB</u>		<u>50MG/25ML (2MG/ML)</u>	<u>A208657 002</u>	May 15, 2017
<u>AB</u>	!	<u>20MG/10ML (2MG/ML)</u>	<u>A203263 001</u>	Feb 04, 2013
<u>AB</u>	!	<u>50MG/25ML (2MG/ML)</u>	<u>A203263 002</u>	Feb 04, 2013

PRESCRIPTION DRUG PRODUCT LIST

DOXYCYCLINE

CAPSULE;ORAL

DOXYCYCLINE

AB	ACP NIMBLE	<u>EQ 50MG BASE</u>	<u>A204446 001</u>	May 28, 2015
AB		<u>EQ 75MG BASE</u>	<u>A204446 002</u>	May 28, 2015
AB		<u>EQ 100MG BASE</u>	<u>A204446 003</u>	May 28, 2015
AB	ALEMBIC PHARMS LTD	<u>EQ 75MG BASE</u>	<u>A209165 001</u>	Jul 28, 2017
AB		<u>EQ 100MG BASE</u>	<u>A209165 002</u>	Jul 28, 2017
AB	IMPAX LABS INC	<u>EQ 150MG BASE</u>	<u>A200065 001</u>	Feb 17, 2011
AB	LUPIN LTD	<u>EQ 50MG BASE</u>	<u>A204234 001</u>	Mar 05, 2014
AB		<u>EQ 75MG BASE</u>	<u>A204234 002</u>	Mar 05, 2014
AB		<u>EQ 100MG BASE</u>	<u>A204234 003</u>	Mar 05, 2014
AB	MAYNE PHARMA INC	<u>EQ 50MG BASE</u>	<u>A209396 001</u>	Sep 29, 2017
AB		<u>EQ 75MG BASE</u>	<u>A209396 002</u>	Sep 29, 2017
AB		<u>EQ 100MG BASE</u>	<u>A209396 003</u>	Sep 29, 2017
AB	PAR PHARM	<u>EQ 50MG BASE</u>	<u>A065055 001</u>	Dec 01, 2000
AB		<u>EQ 100MG BASE</u>	<u>A065055 002</u>	Dec 01, 2000
AB	!	<u>EQ 150MG BASE</u>	<u>A065055 003</u>	Jul 15, 2005
AB	SUN PHARM INDS LTD	<u>EQ 50MG BASE</u>	<u>A065053 001</u>	Nov 22, 2000
AB		<u>EQ 75MG BASE</u>	<u>A065053 003</u>	Sep 10, 2003
AB	!	<u>EQ 100MG BASE</u>	<u>A065053 002</u>	Nov 22, 2000
AB	ZYDUS PHARMS	<u>EQ 50MG BASE</u>	<u>A205115 001</u>	Feb 18, 2016
AB		<u>EQ 75MG BASE</u>	<u>A205115 002</u>	Feb 18, 2016
AB		<u>EQ 100MG BASE</u>	<u>A205115 003</u>	Feb 18, 2016

ORACEA

+! GALDERMA LABS LP 40MG

N050805 001 May 26, 2006

FOR SUSPENSION;ORAL

DOXYCYCLINE

AB	CHARTWELL LIFE SCI	<u>EQ 25MG BASE/5ML</u>	<u>A065454 001</u>	Jul 16, 2008
AB	LUPIN LTD	<u>EQ 25MG BASE/5ML</u>	<u>A201678 001</u>	Mar 18, 2013

VIBRAMYCIN

AB	+! PFIZER	<u>EQ 25MG BASE/5ML</u>	<u>N050006 001</u>	
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TABLET;ORAL

DOXYCYCLINE

AB	HERITAGE PHARMS INC	<u>EQ 50MG BASE</u>	<u>A091605 001</u>	Dec 20, 2011
AB		<u>EQ 75MG BASE</u>	<u>A091605 002</u>	Dec 20, 2011
AB		<u>EQ 100MG BASE</u>	<u>A091605 003</u>	Dec 20, 2011
AB	!	<u>EQ 150MG BASE</u>	<u>A091605 004</u>	Dec 20, 2011
AB	LANNETT CO INC	<u>EQ 50MG BASE</u>	<u>A065285 001</u>	Dec 08, 2005
AB		<u>EQ 75MG BASE</u>	<u>A065285 003</u>	Jul 30, 2008
AB		<u>EQ 100MG BASE</u>	<u>A065285 002</u>	Dec 08, 2005
AB		<u>EQ 150MG BASE</u>	<u>A065285 004</u>	Jul 30, 2008
AB	SUN PHARM INDS LTD	<u>EQ 50MG BASE</u>	<u>A065356 001</u>	May 31, 2006
AB		<u>EQ 75MG BASE</u>	<u>A065356 002</u>	May 31, 2006
AB		<u>EQ 100MG BASE</u>	<u>A065356 003</u>	May 31, 2006
AB		<u>EQ 150MG BASE</u>	<u>A065356 004</u>	Jul 29, 2010
AB	ZYDUS PHARMS	<u>EQ 50MG BASE</u>	<u>A209582 001</u>	Sep 28, 2017
AB		<u>EQ 75MG BASE</u>	<u>A209582 002</u>	Sep 28, 2017
AB		<u>EQ 100MG BASE</u>	<u>A209582 003</u>	Sep 28, 2017
AB		<u>EQ 150MG BASE</u>	<u>A209582 004</u>	Sep 28, 2017

DOXYCYCLINE CALCIUM

SUSPENSION;ORAL

VIBRAMYCIN

+! PFIZER EQ 50MG BASE/5ML

N050480 001

DOXYCYCLINE HYCLATE

CAPSULE;ORAL

DOXYCYCLINE HYCLATE

AB	ACTAVIS LABS FL INC	<u>EQ 50MG BASE</u>	<u>A062031 002</u>	Oct 13, 1982
AB		<u>EQ 100MG BASE</u>	<u>A062031 001</u>	
AB	ALEMBIC PHARMS LTD	<u>EQ 50MG BASE</u>	<u>A210527 001</u>	Jun 13, 2018
AB		<u>EQ 100MG BASE</u>	<u>A210527 002</u>	Jun 13, 2018
AB	AMNEAL PHARMS	<u>EQ 100MG BASE</u>	<u>A207289 001</u>	Jun 27, 2016
AB	CHANGZHOU PHARM	<u>EQ 100MG BASE</u>	<u>A209402 001</u>	Oct 07, 2019
AB	CHARTWELL LIFE SCI	<u>EQ 50MG BASE</u>	<u>A062500 001</u>	Sep 11, 1984
AB		<u>EQ 100MG BASE</u>	<u>A062500 002</u>	Sep 11, 1984
AB	HIKMA INTL PHARMS	<u>EQ 50MG BASE</u>	<u>A062396 002</u>	Nov 07, 1984
AB		<u>EQ 100MG BASE</u>	<u>A062396 001</u>	May 07, 1984
AB	SUN PHARM INDUSTRIES	<u>EQ 50MG BASE</u>	<u>A062676 002</u>	Jul 10, 1986
AB		<u>EQ 100MG BASE</u>	<u>A062676 001</u>	Jul 10, 1986
AB	ZYDUS PHARMS	<u>EQ 50MG BASE</u>	<u>A207774 001</u>	May 31, 2018
AB		<u>EQ 100MG BASE</u>	<u>A207774 002</u>	May 31, 2018

PRESCRIPTION DRUG PRODUCT LIST

DOXYCYCLINE HYCLATE

CAPSULE; ORAL

VIBRAMYCIN

AB	+	PFIZER	<u>EQ 100MG BASE</u>	<u>N050007 002</u>	
INJECTABLE; INJECTION					
<u>DOXY 100</u>					
AP	!	FRESENIUS KABI USA	<u>EQ 100MG BASE/VIAL</u>	<u>A062475 001</u>	Dec 09, 1983
<u>DOXY 200</u>					
AP	!	FRESENIUS KABI USA	<u>EQ 200MG BASE/VIAL</u>	<u>A062475 002</u>	Dec 09, 1983
<u>DOXYCYCLINE</u>					
AP		MYLAN LABS LTD	<u>EQ 100MG BASE/VIAL</u>	<u>A091406 001</u>	Aug 21, 2012
AP	!	WEST-WARD PHARMS INT	<u>EQ 100MG BASE/VIAL</u>	<u>A062569 001</u>	Mar 09, 1988
AP		ZYDUS PHARMS	<u>EQ 100MG BASE/VIAL</u>	<u>A207757 001</u>	Sep 28, 2017
AP			<u>EQ 200MG BASE/VIAL</u>	<u>A207757 002</u>	Sep 28, 2017

SYSTEM, EXTENDED RELEASE; PERIODONTAL

ATRIDOX

+! TOLMAR

50MG

N050751 001 Sep 03, 1998

TABLET; ORAL

ACTICLATE

AB	+	ALMIRALL	<u>EQ 75MG BASE</u>	<u>N205931 001</u>	Jul 25, 2014
AB	+		<u>EQ 150MG BASE</u>	<u>N205931 002</u>	Jul 25, 2014

DOXYCYCLINE HYCLATE

AB		ACTAVIS LABS FL INC	<u>EQ 100MG BASE</u>	<u>A062421 001</u>	Feb 02, 1983
AB		ALEMBIC PHARMS LTD	<u>EQ 20MG BASE</u>	<u>A210537 001</u>	Mar 03, 2020
AB		AMNEAL PHARMS CO	<u>EQ 75MG BASE</u>	<u>A209372 001</u>	Oct 06, 2017
AB			<u>EQ 150MG BASE</u>	<u>A209372 002</u>	Oct 06, 2017
AB		APOTEX	<u>EQ 75MG BASE</u>	<u>A209243 001</u>	Apr 15, 2019
AB			<u>EQ 150MG BASE</u>	<u>A209243 002</u>	Apr 15, 2019
AB		CADILA	<u>EQ 100MG BASE</u>	<u>A207773 001</u>	Oct 30, 2017
AB		CARIBE HOLDINGS	<u>EQ 100MG BASE</u>	<u>A062269 002</u>	Nov 08, 1982
AB		CHANGZHOU PHARM	<u>EQ 100MG BASE</u>	<u>A211343 001</u>	Oct 09, 2019
AB		CHARTWELL LIFE SCI	<u>EQ 100MG BASE</u>	<u>A062505 001</u>	Sep 11, 1984
AB		EMCURE PHARMS LTD	<u>EQ 100MG BASE</u>	<u>A209969 001</u>	Nov 09, 2018
AB		EPIC PHARMA LLC	<u>EQ 20MG BASE</u>	<u>A065182 001</u>	May 13, 2005
AB		HERITAGE PHARMA	<u>EQ 20MG BASE</u>	<u>A065163 001</u>	May 13, 2005
AB	!	HIKMA INTL PHARMS	<u>EQ 100MG BASE</u>	<u>A065095 001</u>	Jul 02, 2003
AB	!	LANNETT CO INC	<u>EQ 20MG BASE</u>	<u>A065277 001</u>	Nov 10, 2005
AB		LARKEN LABS	<u>EQ 20MG BASE</u>	<u>A065287 001</u>	Feb 28, 2006
AB		LUPIN LTD	<u>EQ 75MG BASE</u>	<u>A208818 001</u>	Sep 27, 2017
AB			<u>EQ 150MG BASE</u>	<u>A208818 002</u>	Sep 27, 2017
AB		MAYNE PHARMA INC	<u>EQ 75MG BASE</u>	<u>A208765 001</u>	Jun 14, 2017
AB			<u>EQ 150MG BASE</u>	<u>A208765 002</u>	Jun 14, 2017
AB		MYLAN	<u>EQ 100MG BASE</u>	<u>A062432 001</u>	Feb 15, 1983
AB		NOVEL LABS INC	<u>EQ 100MG BASE</u>	<u>A207558 001</u>	Sep 06, 2017
AB		OAKLOCK LLC	<u>EQ 100MG BASE</u>	<u>A210664 001</u>	Mar 16, 2020
AB		SUN PHARM INDUSTRIES	<u>EQ 20MG BASE</u>	<u>A065134 001</u>	May 13, 2005
AB			<u>EQ 100MG BASE</u>	<u>A062677 001</u>	Jul 10, 1986
		CARIBE HOLDINGS	EQ 50MG BASE	A062269 003	Oct 05, 1983

TABLET, DELAYED RELEASE; ORAL

DORYX

AB	+	MAYNE PHARMA	<u>EQ 50MG BASE</u>	<u>N050795 006</u>	Dec 19, 2014
AB	+		<u>EQ 75MG BASE</u>	<u>N050795 001</u>	May 06, 2005
AB	+		<u>EQ 100MG BASE</u>	<u>N050795 002</u>	May 06, 2005
AB	+		<u>EQ 150MG BASE</u>	<u>N050795 003</u>	Jun 20, 2008
AB	+		<u>EQ 200MG BASE</u>	<u>N050795 005</u>	Apr 11, 2013

DOXYCYCLINE HYCLATE

AB		ACTAVIS ELIZABETH	<u>EQ 50MG BASE</u>	<u>A090134 003</u>	May 22, 2018
AB			<u>EQ 75MG BASE</u>	<u>A090134 001</u>	Dec 14, 2011
AB			<u>EQ 100MG BASE</u>	<u>A090134 002</u>	Dec 14, 2011
AB			<u>EQ 200MG BASE</u>	<u>A090134 004</u>	May 22, 2018
AB		HERITAGE PHARMS INC	<u>EQ 75MG BASE</u>	<u>A200856 001</u>	Apr 30, 2013
AB			<u>EQ 100MG BASE</u>	<u>A200856 002</u>	Apr 30, 2013
AB			<u>EQ 150MG BASE</u>	<u>A200856 003</u>	Apr 30, 2013
AB			<u>EQ 200MG BASE</u>	<u>A200856 004</u>	Nov 13, 2018
AB		MYLAN	<u>EQ 50MG BASE</u>	<u>A090431 003</u>	May 23, 2016
AB		PRINSTON INC	<u>EQ 50MG BASE</u>	<u>A207494 003</u>	Feb 19, 2019
AB			<u>EQ 150MG BASE</u>	<u>A207494 001</u>	Nov 15, 2016
AB			<u>EQ 200MG BASE</u>	<u>A207494 002</u>	Nov 15, 2016

DORYX MPC

+! MAYNE PHARMA

EQ 120MG BASE

N050795 008 May 20, 2016

PRESCRIPTION DRUG PRODUCT LIST

DOXYLAMINE SUCCINATE; PYRIDOXINE HYDROCHLORIDE

TABLET, DELAYED RELEASE; ORAL

DICLEGIS

AB	+ !	DUCHESNAY	10MG;10MG	N021876 001	Apr 08, 2013
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DOXYLAMINE SUCCINATE AND PYRIDOXINE HYDROCHLORIDE

AB		ACTAVIS LABS FL INC	10MG;10MG	A205811 001	Aug 19, 2016
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AB		PAR PHARM INC	10MG;10MG	A208518 001	Dec 06, 2017
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TABLET, EXTENDED RELEASE; ORAL

BONJESTA

+ !	DUCHESNAY	20MG;20MG	N209661 001	Nov 07, 2016
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DRONABINOL

CAPSULE; ORAL

DRONABINOL

AB		AKORN INC	2.5MG	A079217 001	Jun 20, 2014
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AB			5MG	A079217 002	Jun 20, 2014
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AB			10MG	A079217 003	Jun 20, 2014
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AB		ASCENT PHARMS INC	2.5MG	A207421 001	Feb 07, 2020
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AB			5MG	A207421 002	Feb 07, 2020
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AB			10MG	A207421 003	Feb 07, 2020
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AB		LANNETT CO INC	2.5MG	A201463 001	May 18, 2018
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AB			5MG	A201463 002	May 18, 2018
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AB			10MG	A201463 003	May 18, 2018
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AB		SVC PHARMA	2.5MG	A078292 001	Jun 27, 2008
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AB			5MG	A078292 002	Jun 27, 2008
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AB			10MG	A078292 003	Jun 27, 2008
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MARINOL

AB	+	ALKEM LABS LTD	2.5MG	N018651 001	May 31, 1985
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AB	+ !		5MG	N018651 002	May 31, 1985
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AB	+		10MG	N018651 003	May 31, 1985
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SOLUTION; ORAL

SYNDROS

+ !	BENUVIA	5MG/ML	N205525 001	Mar 23, 2017
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DRONEDARONE HYDROCHLORIDE

TABLET; ORAL

MULTAQ

+ !	SANOFI AVENTIS US	EQ 400MG BASE	N022425 001	Jul 01, 2009
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DROPERIDOL

INJECTABLE; INJECTION

DROPERIDOL

AP		AM REGENT	2.5MG/ML	A072123 001	Oct 24, 1988
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AP		EUROHLTH INTL SARL	2.5MG/ML	A208197 001	Dec 14, 2017
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AP		HOSPIRA	2.5MG/ML	A071981 001	Feb 29, 1988
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INAPSINE

AP	+ !	AKORN INC	2.5MG/ML	N016796 001	
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DROSPIRENONE

TABLET; ORAL

SLYND

+ !	EXELTIS USA INC	4MG	N211367 001	May 23, 2019
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DROSPIRENONE; ESTRADIOL

TABLET; ORAL

ANGELIQ

+	BAYER HLTHCARE	0.25MG; 0.5MG	N021355 001	Feb 29, 2012
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+ !		0.5MG; 1MG	N021355 002	Sep 28, 2005
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DROSPIRENONE; ETHINYL ESTRADIOL

TABLET; ORAL

DROSPIRENONE AND ETHINYL ESTRADIOL

AB		ALLIED	3MG;0.02MG	A203291 001	Jul 18, 2017
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AB		BARR	3MG;0.02MG	A078515 001	Mar 30, 2009
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AB		GLENMARK PHARMS LTD	3MG;0.02MG	A204296 001	Aug 17, 2015
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AB		HETERO LABS LTD	3MG;0.02MG	A211944 001	Mar 22, 2019
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AB		JUBILANT CADISTA	3MG;0.02MG	A209423 001	Dec 22, 2017
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AB		MYLAN LABS LTD	3MG;0.02MG	A202594 001	Oct 22, 2015
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AB		WATSON LABS	3MG;0.02MG	A078833 001	Nov 28, 2011
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KYRA

AB		SUN PHARM	3MG;0.02MG	A202318 001	Jul 23, 2019
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LO-ZUMANDIMINE

AB		AUROBINDO PHARMA LTD	3MG;0.02MG	A209632 001	Feb 27, 2018
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PRESCRIPTION DRUG PRODUCT LISTDROSPIRENONE; ETHINYL ESTRADIOL

TABLET; ORAL

LORYNA

AB	XIROMED	3MG;0.02MG	A079221 001	Mar 28, 2011
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MELAMISA

AB	NOVAST LABS	3MG;0.02MG	A202016 001	Jan 26, 2016
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NIKKI

AB	LUPIN LTD	3MG;0.02MG	A201661 001	May 27, 2014
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YAZ

AB	+! BAYER HLTHCARE	3MG;0.02MG	N021676 001	Mar 16, 2006
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TABLET; ORAL-28

DROSPIRENONE AND ETHINYL ESTRADIOL

AB	ACCORD HLTHCARE	3MG;0.03MG	A207245 001	Nov 22, 2016
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AB	APOTEX	3MG;0.03MG	A205876 001	Sep 21, 2016
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AB	BARR	3MG;0.03MG	A077527 001	May 09, 2008
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AB	GLENMARK PHARMS LTD	3MG;0.03MG	A204848 001	Mar 25, 2016
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AB	HETERO LABS LTD	3MG;0.03MG	A213034 001	Jan 24, 2020
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AB	JUBILANT CADISTA	3MG;0.03MG	A210017 001	Sep 10, 2018
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AB	LUPIN LTD	3MG;0.03MG	A201663 001	Dec 18, 2012
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AB	MAYNE PHARMA	3MG;0.03MG	A090081 001	Sep 07, 2010
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AB	MYLAN LABS LTD	3MG;0.03MG	A202131 001	May 04, 2015
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KEMEYA

AB	SUN PHARM	3MG;0.03MG	A202138 001	Mar 13, 2019
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SYEDA

AB	XIROMED	3MG;0.03MG	A090114 001	Mar 28, 2011
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YAEIA

AB	NOVAST LABS	3MG;0.03MG	A202015 001	Nov 19, 2014
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YASMIN

AB	+! BAYER HLTHCARE	3MG;0.03MG	N021098 001	May 11, 2001
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ZUMANDIMINE

AB	AUROBINDO PHARMA LTD	3MG;0.03MG	A209407 001	Mar 26, 2018
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DROSPIRENONE; ETHINYL ESTRADIOL; LEVOMEFOLATE CALCIUM

TABLET; ORAL

BEYAZ

AB	BAYER HLTHCARE	3MG,N/A;0.02MG,N/A;0.451MG,0.451MG	N022532 001	Sep 24, 2010
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DROSPIRENONE, ETHINYL ESTRADIOL AND LEVOMEFOLATE CALCIUM

AB	LUPIN LTD	3MG,N/A;0.02MG,N/A;0.451MG,0.451MG	A205947 001	Jun 13, 2018
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AB	WATSON LABS INC	3MG,N/A;0.02MG,N/A;0.451MG,0.451MG	A203593 001	Oct 11, 2016
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AB		3MG,N/A;0.03MG,N/A;0.451MG,0.451MG	A203594 001	Oct 11, 2016
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SAFYRAL

AB	+! BAYER HLTHCARE	3MG,N/A;0.03MG,N/A;0.451MG,0.451MG	N022574 001	Dec 16, 2010
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TYDEMY

AB	LUPIN LTD	3MG,N/A;0.03MG,N/A;0.451MG,0.451MG	A205948 001	Dec 12, 2017
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DROXIDOPA

CAPSULE; ORAL

NORTHERA

+	LUNDBECK NA LTD	100MG	N203202 001	Feb 18, 2014
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+		200MG	N203202 002	Feb 18, 2014
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+!		300MG	N203202 003	Feb 18, 2014
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DULOXETINE HYDROCHLORIDE

CAPSULE, DELAYED REL PELLETS; ORAL

CYMBALTA

AB	+	LILLY	EQ 20MG BASE	N021427 001	Aug 03, 2004
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AB	+		EQ 30MG BASE	N021427 002	Aug 03, 2004
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AB	+!		EQ 60MG BASE	N021427 004	Aug 03, 2004
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DULOXETINE HYDROCHLORIDE

AB	ACTAVIS ELIZABETH	EQ 20MG BASE	A090776 001	Dec 17, 2013
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AB		EQ 30MG BASE	A090776 002	Dec 17, 2013
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AB		EQ 60MG BASE	A090776 003	Dec 17, 2013
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AB	AJANTA PHARMA LTD	EQ 20MG BASE	A208706 001	Jan 06, 2017
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AB		EQ 30MG BASE	A208706 002	Jan 06, 2017
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AB		EQ 40MG BASE	A208706 004	Mar 11, 2019
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AB		EQ 60MG BASE	A208706 003	Jan 06, 2017
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AB	ALEMBIC PHARMS LTD	EQ 20MG BASE	A202949 001	Jun 09, 2014
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AB		EQ 30MG BASE	A202949 002	Jun 09, 2014
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AB		EQ 60MG BASE	A202949 003	Jun 09, 2014
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AB	ALKEM LABS LTD	EQ 20MG BASE	A203197 001	Aug 26, 2015
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AB		EQ 30MG BASE	A203197 002	Aug 26, 2015
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AB		EQ 60MG BASE	A203197 003	Aug 26, 2015
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AB	ANCHEN PHARMS	EQ 20MG BASE	A090780 001	Oct 28, 2015
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PRESCRIPTION DRUG PRODUCT LIST

DULOXETINE HYDROCHLORIDE

CAPSULE, DELAYED REL PELLETS;ORAL

DULOXETINE HYDROCHLORIDE

<u>AB</u>		<u>EQ 30MG BASE</u>	<u>A090780 002</u>	Oct 28, 2015
<u>AB</u>		<u>EQ 60MG BASE</u>	<u>A090780 003</u>	Oct 28, 2015
<u>AB</u>	APOTEX	<u>EQ 20MG BASE</u>	<u>A202045 001</u>	Jun 11, 2014
<u>AB</u>		<u>EQ 30MG BASE</u>	<u>A202045 002</u>	Jun 11, 2014
<u>AB</u>		<u>EQ 60MG BASE</u>	<u>A202045 003</u>	Jun 11, 2014
<u>AB</u>	AUROBINDO PHARMA LTD	<u>EQ 20MG BASE</u>	<u>A090778 001</u>	Dec 11, 2013
<u>AB</u>		<u>EQ 30MG BASE</u>	<u>A090778 002</u>	Dec 11, 2013
<u>AB</u>		<u>EQ 60MG BASE</u>	<u>A090778 003</u>	Dec 11, 2013
<u>AB</u>	BRECKENRIDGE	<u>EQ 20MG BASE</u>	<u>A203088 001</u>	Jun 11, 2014
<u>AB</u>		<u>EQ 30MG BASE</u>	<u>A203088 002</u>	Jun 11, 2014
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A203088 004</u>	May 18, 2018
<u>AB</u>		<u>EQ 60MG BASE</u>	<u>A203088 003</u>	Jun 11, 2014
<u>AB</u>	CSPC OUYI	<u>EQ 20MG BASE</u>	<u>A211310 001</u>	Oct 16, 2018
<u>AB</u>		<u>EQ 30MG BASE</u>	<u>A211310 002</u>	Oct 16, 2018
<u>AB</u>		<u>EQ 60MG BASE</u>	<u>A211310 003</u>	Oct 16, 2018
<u>AB</u>	HETERO LABS LTD III	<u>EQ 20MG BASE</u>	<u>A204343 001</u>	Aug 03, 2016
<u>AB</u>		<u>EQ 30MG BASE</u>	<u>A204343 002</u>	Aug 03, 2016
<u>AB</u>		<u>EQ 60MG BASE</u>	<u>A204343 003</u>	Aug 03, 2016
<u>AB</u>	INVENTIA	<u>EQ 20MG BASE</u>	<u>A202336 001</u>	Oct 28, 2015
<u>AB</u>		<u>EQ 30MG BASE</u>	<u>A202336 002</u>	Oct 28, 2015
<u>AB</u>		<u>EQ 60MG BASE</u>	<u>A202336 003</u>	Oct 28, 2015
<u>AB</u>	LUPIN LTD	<u>EQ 20MG BASE</u>	<u>A090694 001</u>	Dec 11, 2013
<u>AB</u>		<u>EQ 30MG BASE</u>	<u>A090694 002</u>	Dec 11, 2013
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A090694 003</u>	Dec 11, 2013
<u>AB</u>		<u>EQ 60MG BASE</u>	<u>A090694 004</u>	Dec 11, 2013
<u>AB</u>	MACLEODS PHARMS LTD	<u>EQ 20MG BASE</u>	<u>A204815 001</u>	Mar 23, 2017
<u>AB</u>		<u>EQ 30MG BASE</u>	<u>A204815 002</u>	Mar 23, 2017
<u>AB</u>		<u>EQ 60MG BASE</u>	<u>A204815 003</u>	Mar 23, 2017
<u>AB</u>	MARKSANS PHARMA	<u>EQ 20MG BASE</u>	<u>A090723 001</u>	Dec 11, 2013
<u>AB</u>		<u>EQ 30MG BASE</u>	<u>A090723 002</u>	Dec 11, 2013
<u>AB</u>		<u>EQ 60MG BASE</u>	<u>A090723 003</u>	Dec 11, 2013
<u>AB</u>	PRINSTON INC	<u>EQ 20MG BASE</u>	<u>A206653 001</u>	May 18, 2017
<u>AB</u>		<u>EQ 30MG BASE</u>	<u>A206653 002</u>	May 18, 2017
<u>AB</u>		<u>EQ 60MG BASE</u>	<u>A206653 003</u>	May 18, 2017
<u>AB</u>	QINGDAO BAHEAL PHARM	<u>EQ 20MG BASE</u>	<u>A210599 001</u>	Apr 17, 2019
<u>AB</u>		<u>EQ 30MG BASE</u>	<u>A210599 002</u>	Apr 17, 2019
<u>AB</u>		<u>EQ 60MG BASE</u>	<u>A210599 003</u>	Apr 17, 2019
<u>AB</u>	SUN PHARM	<u>EQ 20MG BASE</u>	<u>A090745 001</u>	Dec 11, 2013
<u>AB</u>		<u>EQ 30MG BASE</u>	<u>A090745 002</u>	Dec 11, 2013
<u>AB</u>		<u>EQ 60MG BASE</u>	<u>A090745 003</u>	Dec 11, 2013
<u>AB</u>	TORRENT	<u>EQ 20MG BASE</u>	<u>A090774 001</u>	Dec 11, 2013
<u>AB</u>		<u>EQ 30MG BASE</u>	<u>A090774 002</u>	Dec 11, 2013
<u>AB</u>		<u>EQ 60MG BASE</u>	<u>A090774 003</u>	Dec 11, 2013
<u>AB</u>	YAOPHARMA CO LTD	<u>EQ 20MG BASE</u>	<u>A207219 001</u>	Aug 16, 2019
<u>AB</u>		<u>EQ 30MG BASE</u>	<u>A207219 002</u>	Aug 16, 2019
<u>AB</u>		<u>EQ 60MG BASE</u>	<u>A207219 003</u>	Aug 16, 2019
<u>AB</u>	ZYDUS HLTHCARE	<u>EQ 20MG BASE</u>	<u>A090739 001</u>	Jan 08, 2014
<u>AB</u>		<u>EQ 30MG BASE</u>	<u>A090739 002</u>	Jan 08, 2014
<u>AB</u>		<u>EQ 60MG BASE</u>	<u>A090739 003</u>	Jan 08, 2014
<u>AB</u>	ZYDUS PHARMS	<u>EQ 20MG BASE</u>	<u>A090728 001</u>	Jan 08, 2014
<u>AB</u>		<u>EQ 30MG BASE</u>	<u>A090728 002</u>	Jan 08, 2014
<u>AB</u>		<u>EQ 60MG BASE</u>	<u>A090728 003</u>	Jan 08, 2014

CAPSULE, DELAYED RELEASE;ORAL

DRIZALMA SPRINKLE

+	SUN PHARMA GLOBAL	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

DUTASTERIDE

CAPSULE;ORAL

AVODART

<u>AB</u>	+	GLAXOSMITHKLINE	<u>0.5MG</u>	<u>N021319 001</u>	Nov 20, 2001
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DUTASTERIDE

<u>AB</u>		AMNEAL PHARMS	<u>0.5MG</u>	<u>A203118 001</u>	Nov 20, 2015
<u>AB</u>		ASCENT PHARMS INC	<u>0.5MG</u>	<u>A206574 001</u>	Oct 21, 2016
<u>AB</u>		AUROLIFE PHARMA LLC	<u>0.5MG</u>	<u>A202660 001</u>	Nov 20, 2015
<u>AB</u>		BARR	<u>0.5MG</u>	<u>A090095 001</u>	Dec 21, 2010

PRESCRIPTION DRUG PRODUCT LISTDUTASTERIDE

CAPSULE; ORAL

DUTASTERIDE

AB	BIONPHARMA INC	0.5MG	A200899 001	Nov 20, 2015
AB	CADILA	0.5MG	A204373 001	Oct 04, 2017
AB	HERITAGE PHARMS INC	0.5MG	A207935 001	Oct 13, 2017
AB	HUMANWELL PURACAP	0.5MG	A209909 001	Nov 21, 2017
AB	INTERGEL PHARMS INC	0.5MG	A206373 001	Mar 17, 2016
AB	MARKSANS PHARMA	0.5MG	A204376 001	Apr 07, 2017
AB	STRIDES PHARMA	0.5MG	A204262 001	Nov 20, 2015
AB	VINTAGE PHARMS LLC	0.5MG	A202421 001	Nov 20, 2015

DUTASTERIDE; TAMSULOSIN HYDROCHLORIDE

CAPSULE; ORAL

DUTASTERIDE AND TAMSULOSIN HYDROCHLORIDE

AB	ACTAVIS LABS FL INC	0.5MG; 0.4MG	A202975 001	Nov 20, 2015
AB	ANCHEN PHARMS	0.5MG; 0.4MG	A202509 001	Feb 26, 2014
AB	ZYDUS PHARMS	0.5MG; 0.4MG	A207769 001	May 24, 2018
JALYN				
AB	+ GLAXOSMITHKLINE	0.5MG; 0.4MG	N022460 001	Jun 14, 2010

DUVELISIB

CAPSULE; ORAL

COPIKTRA

+	VERASTEM INC	15MG	N211155 001	Sep 24, 2018
+	!	25MG	N211155 002	Sep 24, 2018

DYCLONINE HYDROCHLORIDE

SOLUTION; TOPICAL

DYCLOPRO

!	NOVOCOL INC	0.5%	A200480 001	Nov 20, 2018
!		1%	A200480 002	Nov 20, 2018

ECHOTHIOPHATE IODIDE

FOR SOLUTION; OPHTHALMIC

PHOSPHOLINE IODIDE

+	!	WYETH PHARMS	0.125%	N011963 001
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ECONAZOLE NITRATE

AEROSOL, FOAM; TOPICAL

ECOZA

+	!	GLENMARK	1%	N205175 001	Oct 24, 2013
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CREAM; TOPICAL

ECONAZOLE NITRATE

AB	MYLAN	1%	A210364 001	Apr 18, 2018
AB	! PERRIGO ISRAEL	1%	A076479 001	Jun 23, 2004
AB	TARO	1%	A076005 001	Nov 26, 2002
AB	TELIGENT PHARMA INC	1%	A076574 001	Dec 17, 2004

SPECTAZOLE

AB	+	ALVOGEN	1%	N018751 001	Dec 23, 1982
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EDARAVONE

SOLUTION; INTRAVENOUS

RADICAVA

+	!	MITSUBISHI TANABE	30MG/100ML (0.3MG/ML)	N209176 001	May 05, 2017
+	!		60MG/100ML (0.6MG/ML)	N209176 002	Nov 15, 2018

EDETATE CALCIUM DISODIUM

INJECTABLE; INJECTION

CALCIUM DISODIUM VERSENATE

+	!	MEDICIS	200MG/ML	N008922 001
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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

AB	AUROBINDO PHARMA LTD	50MG	A078064 001	Dec 15, 2017	
AB		200MG	A078064 003	Dec 15, 2017	
SUSTIVA					
AB	+	BRISTOL MYERS	50MG	N020972 001	Sep 17, 1998

PRESCRIPTION DRUG PRODUCT LIST

EFAVIRENZ

CAPSULE; ORAL

SUSTIVA

SQUIBB

AB	+		200MG	N020972 003	Sep 17, 1998
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EFAVIRENZ

AUROBINDO PHARMA
LTD

100MG

A078064 002 Dec 15, 2017

TABLET; ORAL

EFAVIRENZ

AB		AUROBINDO PHARMA	600MG	A077673 001	Sep 21, 2018
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LTD

AB			600MG	A205322 001	Aug 30, 2018
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AB		CIPLA	600MG	A204766 001	Jun 15, 2018
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AB		HETERO LABS LTD III	600MG	A078886 001	Apr 27, 2018
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AB		MYLAN	600MG	A091471 001	Feb 17, 2016
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AB		STRIDES PHARMA	600MG	A204869 001	Mar 12, 2018
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SUSTIVA

AB	+	BRISTOL MYERS	600MG	N021360 002	Feb 01, 2002
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SQUIBB

EFAVIRENZ; EMTRICITABINE; TENOFOVIR DISOPROXIL FUMARATE

TABLET; ORAL

ATRIPLA

AB	+	GILEAD SCIENCES	600MG; 200MG; 300MG	N021937 001	Jul 12, 2006
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EFAVIRENZ, EMTRICITABINE, AND TENOFOVIR DISOPROXIL FUMARATE

AB		AUROBINDO PHARMA	600MG; 200MG; 300MG	A203041 001	Sep 04, 2018
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LTD

AB		CIPLA	600MG; 200MG; 300MG	A206894 001	Jun 03, 2019
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EFAVIRENZ; LAMIVUDINE; TENOFOVIR DISOPROXIL FUMARATE

TABLET; ORAL

SYMFI

+! MYLAN LABS LTD

600MG; 300MG; 300MG

N022142 001 Mar 22, 2018

SYMFI LO

+! MYLAN

400MG; 300MG; 300MG

N208255 001 Feb 05, 2018

EFINACONAZOLE

SOLUTION; TOPICAL

JUBLIA

+! BAUSCH

10%

N203567 001 Jun 06, 2014

EFLORNITHINE HYDROCHLORIDE

CREAM; TOPICAL

VANIQA

+! SKINMEDICA

13.9%

N021145 001 Jul 27, 2000

ELAGOLIX SODIUM

TABLET; ORAL

ORILISSA

+ ABBVIE INC

EQ 150MG BASE

N210450 001 Jul 23, 2018

+!

EQ 200MG BASE

N210450 002 Jul 23, 2018

ELBASVIR; GRAZOPREVIR

TABLET; ORAL

ZEPATIER

+! MERCK SHARP DOHME

50MG; 100MG

N208261 001 Jan 28, 2016

ELETRIPTAN HYDROBROMIDE

TABLET; ORAL

ELETRIPTAN HYDROBROMIDE

AB		AJANTA PHARMA LTD	EQ 20MG BASE	A205186 001	Aug 29, 2017
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AB			EQ 40MG BASE	A205186 002	Aug 29, 2017
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AB		AMNEAL PHARMS CO	EQ 20MG BASE	A206787 001	May 25, 2018
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AB			EQ 40MG BASE	A206787 002	May 25, 2018
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AB		AUROBINDO PHARMA	EQ 20MG BASE	A210708 001	Jan 15, 2019
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LTD

AB			EQ 40MG BASE	A210708 002	Jan 15, 2019
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AB		MYLAN	EQ 20MG BASE	A205152 001	Aug 11, 2017
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AB			EQ 40MG BASE	A205152 002	Aug 11, 2017
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AB		TEVA PHARMS USA	EQ 20MG BASE	A202040 001	Jun 27, 2017
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AB			EQ 40MG BASE	A202040 002	Jun 27, 2017
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AB		ZYDUS PHARMS	EQ 20MG BASE	A206409 001	Jun 16, 2017
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AB			EQ 40MG BASE	A206409 002	Jun 16, 2017
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RELPAK

AB	+	PFIZER IRELAND	EQ 20MG BASE	N021016 001	Dec 26, 2002
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AB	+		EQ 40MG BASE	N021016 002	Dec 26, 2002
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PRESCRIPTION DRUG PRODUCT LISTELEXACAFTOR, IVACAFTOR, TEZACAFTOR; IVACAFTOR

TABLET, TABLET;ORAL

TRIKAFTA (COPACKAGED)

+! VERTEX PHARMS INC 100MG,75MG,50MG;N/A,150MG,N/A N212273 001 Oct 21, 2019

ELIGLUSTAT TARTRATE

CAPSULE;ORAL

CERDELGA

+! GENZYME CORP EQ 84MG BASE N205494 001 Aug 19, 2014

ELTROMBOPAG OLAMINE

FOR SUSPENSION;ORAL

PROMACTA KIT

+ NOVARTIS EQ 12.5MG ACID/PACKET N207027 002 Sep 27, 2018

+! 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

+ ALLERGAN HOLDINGS 75MG N206940 001 May 27, 2015

+! 100MG N206940 002 May 27, 2015

EMPAGLIFLOZIN

TABLET;ORAL

JARDIANCE

+ BOEHRINGER 10MG N204629 001 Aug 01, 2014

+! INGELHEIM 25MG N204629 002 Aug 01, 2014

EMPAGLIFLOZIN; LINAGLIPTIN

TABLET;ORAL

GLYXAMBI

+ BOEHRINGER 10MG;5MG N206073 001 Jan 30, 2015

+! INGELHEIM 25MG;5MG N206073 002 Jan 30, 2015

EMPAGLIFLOZIN; LINAGLIPTIN; METFORMIN HYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL

TRIJARDY XR

+ BOEHRINGER 5MG;2.5MG;1GM N212614 001 Jan 27, 2020

+ INGELHEIM 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 5MG;500MG N206111 001 Aug 26, 2015

+ INGELHEIM 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 5MG;1GM N208658 001 Dec 09, 2016

+ INGELHEIM 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** 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

+! GILEAD SCIENCES INC 200MG;EQ 25MG BASE N208215 001 Apr 04, 2016

EMTRICITABINE; TENOFOVIR DISOPROXIL FUMARATE

TABLET;ORAL

EMTRICITABINE AND TENOFOVIR DISOPROXIL FUMARATE

AB	AUROBINDO PHARMA LTD	200MG;300MG	A090513 001	Jan 26, 2018
AB	TEVA PHARMS USA	200MG;300MG	A090894 001	Jun 08, 2017
AB	ZYDUS PHARMS	200MG;300MG	A212689 001	Feb 28, 2020
TRUVADA				
AB	+! GILEAD	200MG;300MG	N021752 001	Aug 02, 2004
	+	100MG;150MG	N021752 002	Mar 10, 2016
	+	133MG;200MG	N021752 003	Mar 10, 2016
	+	167MG;250MG	N021752 004	Mar 10, 2016

ENALAPRIL MALEATE

SOLUTION;ORAL

EPANED

+! SILVERGATE PHARMS 1MG/ML N208686 001 Sep 20, 2016

TABLET;ORAL

ENALAPRIL MALEATE

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	NOSTRUM LABS 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	SANDOZ INC	2.5MG	A075496 001	Aug 22, 2000
AB		5MG	A075496 002	Aug 22, 2000
AB		10MG	A075459 001	Aug 22, 2000
AB		20MG	A075459 002	Aug 22, 2000
AB	TARO	2.5MG	A075657 001	Jan 23, 2001
AB		5MG	A075657 002	Jan 23, 2001
AB		10MG	A075657 003	Jan 23, 2001
AB		20MG	A075657 004	Jan 23, 2001
AB	WOCKHARDT LTD	2.5MG	A075483 001	Aug 22, 2000
AB		5MG	A075483 002	Aug 22, 2000
AB		10MG	A075483 003	Aug 22, 2000
AB		20MG	A075483 004	Aug 22, 2000

VASOTEC

AB	+ BAUSCH	2.5MG	N018998 005	Jul 26, 1988
AB	+	5MG	N018998 001	Dec 24, 1985
AB	+	10MG	N018998 002	Dec 24, 1985
AB	+!	20MG	N018998 003	Dec 24, 1985

ENALAPRIL MALEATE; HYDROCHLOROTHIAZIDE

TABLET;ORAL

ENALAPRIL MALEATE AND HYDROCHLOROTHIAZIDE

AB	ACP NIMBLE	5MG;12.5MG	A075727 001	Sep 18, 2001
AB		10MG;25MG	A075727 002	Sep 18, 2001
AB	DR REDDYS LABS LTD	5MG;12.5MG	A075909 001	Oct 15, 2001
AB		10MG;25MG	A075909 002	Oct 15, 2001
AB	MYLAN	5MG;12.5MG	A075624 001	Sep 18, 2001
AB		10MG;25MG	A075624 002	Sep 18, 2001
AB	NOSTRUM LABS INC	5MG;12.5MG	A076486 001	Oct 27, 2004
AB		10MG;25MG	A076486 002	Oct 27, 2004
AB	TARO PHARM INDS	5MG;12.5MG	A075788 001	Sep 18, 2001
AB		10MG;25MG	A075788 002	Sep 18, 2001

VASERETIC

AB	+ BAUSCH	5MG;12.5MG	N019221 003	Jul 12, 1995
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PRESCRIPTION DRUG PRODUCT LIST

ENALAPRIL MALEATE; HYDROCHLOROTHIAZIDE

TABLET; ORAL

VASERETIC

AB	+ !		10MG;25MG	N019221	001	Oct 31, 1986
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ENALAPRILAT

INJECTABLE; INJECTION

ENALAPRILAT

AP	!	ATHENEX INC	1.25MG/ML	A075634	001	Aug 22, 2000
AP		DR REDDYS	1.25MG/ML	A075578	001	Aug 22, 2000
AP		HIKMA FARMACEUTICA	1.25MG/ML	A078687	001	Dec 23, 2008
AP	!	HOSPIRA	1.25MG/ML	A075458	001	Aug 22, 2000

ENASIDENIB MESYLATE

TABLET; ORAL

IDHIFA

+		CELGENE CORP	EQ 50MG BASE	N209606	001	Aug 01, 2017
+ !			EQ 100MG BASE	N209606	002	Aug 01, 2017

ENCORAFENIB

CAPSULE; ORAL

BRAFTOVI

+ !		ARRAY BIOPHARMA INC	75MG	N210496	002	Jun 27, 2018
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ENFUVRTIDE

INJECTABLE; SUBCUTANEOUS

FUZEON

+ !		ROCHE	90MG/VIAL	N021481	001	Mar 13, 2003
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ENOXAPARIN SODIUM

INJECTABLE; INTRAVENOUS, SUBCUTANEOUS

ENOXAPARIN SODIUM

AB		AMPHASTAR PHARMS INC	300MG/3ML (100MG/ML)	A208600	001	Mar 14, 2019
AB		SANDOZ INC	300MG/3ML (100MG/ML)	A078660	001	Nov 28, 2011
AB	+	SANOFI AVENTIS US	300MG/3ML (100MG/ML)	N020164	009	Jan 23, 2003

INJECTABLE; SUBCUTANEOUS

ENOXAPARIN SODIUM (PRESERVATIVE FREE)

AP		AMPHASTAR PHARM	30MG/0.3ML (100MG/ML)	A076684	001	Sep 19, 2011
AP			40MG/0.4ML (100MG/ML)	A076684	002	Sep 19, 2011
AP			60MG/0.6ML (100MG/ML)	A076684	003	Sep 19, 2011
AP			80MG/0.8ML (100MG/ML)	A076684	004	Sep 19, 2011
AP			100MG/ML (100MG/ML)	A076684	005	Sep 19, 2011
AP			120MG/0.8ML (150MG/ML)	A076684	006	Sep 19, 2011
AP			150MG/ML (150MG/ML)	A076684	007	Sep 19, 2011
AP		APOTEX INC	30MG/0.3ML (100MG/ML)	A078990	001	Sep 28, 2018
AP			40MG/0.4ML (100MG/ML)	A078990	002	Sep 28, 2018
AP			60MG/0.6ML (100MG/ML)	A078990	003	Sep 28, 2018
AP			80MG/0.8ML (100MG/ML)	A078990	004	Sep 28, 2018
AP			100MG/ML (100MG/ML)	A078990	005	Sep 28, 2018
AP			120MG/0.8ML (150MG/ML)	A078990	006	Sep 28, 2018
AP			150MG/ML (150MG/ML)	A078990	007	Sep 28, 2018
AP		NANJING KING-FRIEND	30MG/0.3ML (100MG/ML)	A206834	001	Nov 29, 2019
AP			40MG/0.4ML (100MG/ML)	A206834	002	Nov 29, 2019
AP			60MG/0.6ML (100MG/ML)	A206834	003	Nov 29, 2019
AP			80MG/0.8ML (100MG/ML)	A206834	004	Nov 29, 2019
AP			100MG/ML (100MG/ML)	A206834	005	Nov 29, 2019
AP			120MG/0.8ML (150MG/ML)	A206834	006	Nov 29, 2019
AP			150MG/ML (150MG/ML)	A206834	007	Nov 29, 2019
AP		SANDOZ	30MG/0.3ML (100MG/ML)	A077857	002	Jul 23, 2010
AP			40MG/0.4ML (100MG/ML)	A077857	003	Jul 23, 2010
AP			60MG/0.6ML (100MG/ML)	A077857	004	Jul 23, 2010
AP			80MG/0.8ML (100MG/ML)	A077857	005	Jul 23, 2010
AP			100MG/ML (100MG/ML)	A077857	001	Jul 23, 2010
AP			120MG/0.8ML (150MG/ML)	A077857	006	Jul 23, 2010
AP			150MG/ML (150MG/ML)	A077857	007	Jul 23, 2010
AP		TEVA	30MG/0.3ML (100MG/ML)	A076726	001	Jun 23, 2014
AP			40MG/0.4ML (100MG/ML)	A076726	002	Jun 23, 2014
AP			60MG/0.6ML (100MG/ML)	A076726	003	Jun 23, 2014
AP			80MG/0.8ML (100MG/ML)	A076726	004	Jun 23, 2014
AP			100MG/ML (100MG/ML)	A076726	005	Jun 23, 2014
AP			120MG/0.8ML (150MG/ML)	A076726	006	Jun 23, 2014
AP			150MG/ML (150MG/ML)	A076726	007	Jun 23, 2014

PRESCRIPTION DRUG PRODUCT LIST

ENOXAPARIN SODIUM

INJECTABLE; SUBCUTANEOUS

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

ENTACAPONE

TABLET; ORAL

COMTAN

<u>AB</u>	+	ORION PHARMA	<u>200MG</u>	<u>N020796 001</u>	Oct 19, 1999
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ENTACAPONE

<u>AB</u>		AJANTA PHARMA LTD	<u>200MG</u>	<u>A205792 001</u>	Aug 31, 2017
<u>AB</u>		AUROBINDO PHARMA LTD	<u>200MG</u>	<u>A203437 001</u>	Jun 19, 2015
<u>AB</u>		MACLEODS PHARMS LTD	<u>200MG</u>	<u>A207210 001</u>	Jun 05, 2017
<u>AB</u>		SUN PHARM	<u>200MG</u>	<u>A090690 001</u>	Jul 16, 2012
<u>AB</u>		SUNSHINE LAKE	<u>200MG</u>	<u>A206669 001</u>	Oct 03, 2018
<u>AB</u>		WOCKHARDT LTD	<u>200MG</u>	<u>A078941 001</u>	Aug 16, 2012

ENTECAVIR

SOLUTION; ORAL

BARACLUE

	+	BRISTOL MYERS SQUIBB	0.05MG/ML	N021798 001	Mar 29, 2005
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TABLET; ORAL

BARACLUE

<u>AB</u>	+	BRISTOL MYERS SQUIBB	<u>0.5MG</u>	<u>N021797 001</u>	Mar 29, 2005
<u>AB</u>	+		<u>1MG</u>	<u>N021797 002</u>	Mar 29, 2005

ENTECAVIR

<u>AB</u>		ACCORD HLTHCARE	<u>0.5MG</u>	<u>A205824 001</u>	Aug 25, 2017
<u>AB</u>			<u>1MG</u>	<u>A205824 002</u>	Aug 25, 2017
<u>AB</u>		AMNEAL PHARMS	<u>0.5MG</u>	<u>A206652 001</u>	Nov 12, 2015
<u>AB</u>			<u>1MG</u>	<u>A206652 002</u>	Nov 12, 2015
<u>AB</u>		AUROBINDO PHARMA LTD	<u>0.5MG</u>	<u>A206217 001</u>	Aug 26, 2015
<u>AB</u>			<u>1MG</u>	<u>A206217 002</u>	Aug 26, 2015
<u>AB</u>		BRECKENRIDGE	<u>0.5MG</u>	<u>A208721 001</u>	Mar 15, 2018
<u>AB</u>			<u>1MG</u>	<u>A208721 002</u>	Mar 15, 2018
<u>AB</u>		BRIGHTGENE	<u>0.5MG</u>	<u>A212126 001</u>	Sep 25, 2019
<u>AB</u>			<u>1MG</u>	<u>A212126 002</u>	Sep 25, 2019
<u>AB</u>		CASI PHARMS INC	<u>0.5MG</u>	<u>A206672 001</u>	May 11, 2017
<u>AB</u>			<u>1MG</u>	<u>A206672 002</u>	May 11, 2017
<u>AB</u>		CIPLA	<u>0.5MG</u>	<u>A206872 001</u>	Dec 06, 2016
<u>AB</u>			<u>1MG</u>	<u>A206872 002</u>	Dec 06, 2016
<u>AB</u>		HETERO LABS LTD V	<u>0.5MG</u>	<u>A205740 001</u>	Aug 21, 2015
<u>AB</u>			<u>1MG</u>	<u>A205740 002</u>	Aug 21, 2015
<u>AB</u>		PRINSTON INC	<u>0.5MG</u>	<u>A208782 001</u>	Oct 10, 2017
<u>AB</u>			<u>1MG</u>	<u>A208782 002</u>	Oct 10, 2017
<u>AB</u>		TEVA PHARMS USA	<u>0.5MG</u>	<u>A202122 001</u>	Aug 26, 2014
<u>AB</u>			<u>1MG</u>	<u>A202122 002</u>	Aug 26, 2014
<u>AB</u>		YAOPHARMA CO LTD	<u>0.5MG</u>	<u>A212201 001</u>	Nov 04, 2019
<u>AB</u>			<u>1MG</u>	<u>A212201 002</u>	Nov 04, 2019
<u>AB</u>		ZYDUS PHARMS	<u>0.5MG</u>	<u>A206745 001</u>	Jun 23, 2017
<u>AB</u>			<u>1MG</u>	<u>A206745 002</u>	Jun 23, 2017

ENTRECTINIB

CAPSULE; ORAL

ROZLYTREK

	+	GENENTECH INC	100MG	N212725 001	Aug 15, 2019
	+		200MG	N212725 002	Aug 15, 2019

ENZALUTAMIDE

CAPSULE; ORAL

XTANDI

	+	ASTELLAS	40MG	N203415 001	Aug 31, 2012
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PRESCRIPTION DRUG PRODUCT LIST

EPHEDRINE SULFATE

SOLUTION; INTRAVENOUS

AKOVAZ

AP	+!	AVADEL LEGACY	50MG/ML (50MG/ML)	N208289 001	Apr 29, 2016
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CORPHEDRA

AP		PAR STERILE PRODUCTS	50MG/ML (50MG/ML)	N208943 001	Jan 27, 2017
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EPHEDRINE SULFATE

AP		AKORN INC	50MG/ML (50MG/ML)	N208609 001	Mar 01, 2017
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AP		AMNEAL	50MG/ML (50MG/ML)	A212932 001	Oct 23, 2019
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AP		SANDOZ INC	50MG/ML (50MG/ML)	A209784 001	Aug 23, 2017
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EPINASTINE HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

ELESTAT

AT	+!	ALLERGAN	0.05%	N021565 001	Oct 16, 2003
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EPINASTINE HYDROCHLORIDE

AT		AKORN	0.05%	A204055 001	May 05, 2017
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AT		APOTEX	0.05%	A090919 001	Oct 31, 2011
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AT		BRECKENRIDGE	0.05%	A090870 001	Mar 14, 2011
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AT		SOMERSET THERAPS LLC	0.05%	A090951 001	Oct 31, 2011
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EPINEPHRINE

INJECTABLE; INTRAMUSCULAR, SUBCUTANEOUS

EPINEPHRINE (AUTOINJECTOR)

AB		TEVA PHARMS USA	0.15MG/DELIVERY	A090589 002	Aug 16, 2018
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AB			0.3MG/DELIVERY	A090589 001	Aug 16, 2018
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EPIPEN

AB	+!	MYLAN SPECIALITY LP	0.3MG/DELIVERY	N019430 001	Dec 22, 1987
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EPIPEN JR.

AB	+!	MYLAN SPECIALITY LP	0.15MG/DELIVERY	N019430 002	Dec 22, 1987
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ADRENACLICK

BX	+!	IMPAX	EQ 0.15MG/DELIVERY	N020800 003	Nov 25, 2009
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BX	+!		EQ 0.3MG/DELIVERY	N020800 004	Nov 25, 2009
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SOLUTION; INTRAMUSCULAR, INTRAVENOUS, SUBCUTANEOUS

ADRENALIN

+!	PAR STERILE PRODUCTS	EQ 1MG BASE/ML (EQ 1MG BASE/ML)	N204200 001	Dec 07, 2012
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+!		EQ 30MG BASE/30ML (EQ 1MG BASE/ML)	N204640 001	Dec 18, 2013
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SOLUTION; INTRAMUSCULAR, SUBCUTANEOUS

AUVI-Q

BX	+!	KALEO INC	EQ 0.15MG/DELIVERY	N201739 002	Aug 10, 2012
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BX	+		EQ 0.3MG/DELIVERY	N201739 001	Aug 10, 2012
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+			EQ 0.1MG/DELIVERY	N201739 003	Nov 17, 2017
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SYMJEPI

+!	ADAMIS PHARMS CORP	0.15MG/0.3ML (0.15MG/0.3ML)	N207534 002	Sep 27, 2018
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+!		0.3MG/0.3ML (0.3MG/0.3ML)	N207534 001	Jun 15, 2017
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SOLUTION; INTRAVENOUS

EPINEPHRINE (COPACKAGED)

+!	HOSPIRA INC	1MG/10ML (0.1MG/ML)	N209359 001	Nov 05, 2019
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SOLUTION; INTRAVENOUS, INTRAOCULAR, INTRAMUSCULAR, SUBCUTANEOUS

EPINEPHRINE

+!	BELCHER	EQ 1MG BASE/ML (EQ 1MG BASE/ML)	N205029 001	Jul 29, 2014
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EPINEPHRINE BITARTRATE; LIDOCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

LIGNOSPAN FORTE

!	DEPROCO	EQ 0.02MG BASE/ML; 2%	A088389 001	Jan 22, 1985
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LIGNOSPAN STANDARD

!	DEPROCO	EQ 0.01MG BASE/ML; 2%	A088390 001	Jan 22, 1985
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EPINEPHRINE BITARTRATE; PRILUCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

CITANEST FORTE DENTAL

AP	+!	DENTSPLY PHARM	0.005MG/ML; 4%	N021383 001	
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PRILOCAINE HYDROCHLORIDE AND EPINEPHRINE BITARTRATE

AP		SEPTODONT INC	0.005MG/ML; 4%	A078959 001	Aug 30, 2011
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PRESCRIPTION DRUG PRODUCT LIST

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EPINEPHRINE; LIDOCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

LIDOCAINE HYDROCHLORIDE AND EPINEPHRINE

<u>AP</u>	HOSPIRA	<u>0.005MG/ML;0.5%</u>	<u>A089635 001</u>	Jun 21, 1988
<u>AP</u>		<u>0.005MG/ML;1.5%</u>	<u>A088571 001</u>	Sep 13, 1985
<u>AP</u>		<u>0.005MG/ML;1.5%</u>	<u>A089645 001</u>	Jun 21, 1988
<u>AP</u>		<u>0.005MG/ML;2%</u>	<u>A089651 001</u>	Jun 21, 1988
<u>AP</u>		<u>0.01MG/ML;1%</u>	<u>A089644 001</u>	Jun 21, 1988
<u>AP</u>	!	<u>0.01MG/ML;2%</u>	<u>A089646 001</u>	Jun 21, 1988
<u>XYLOCAINE W/ EPINEPHRINE</u>				
<u>AP</u>	+!	FRESENIUS KABI USA	<u>0.005MG/ML;0.5%</u>	<u>N006488 012</u>
<u>AP</u>	+!		<u>0.005MG/ML;1.5%</u>	<u>N006488 017</u>
<u>AP</u>	+!		<u>0.005MG/ML;2%</u>	<u>N006488 019</u> Nov 13, 1986
<u>AP</u>	+!		<u>0.01MG/ML;1%</u>	<u>N006488 004</u>
<u>AP</u>	+!		<u>0.02MG/ML;2%</u>	<u>N006488 005</u>
			0.005MG/ML;1%	N006488 018 Nov 13, 1986

EPIRUBICIN HYDROCHLORIDE

INJECTABLE; INJECTION

ELLEENCE

<u>AP</u>	+!	PFIZER INC	<u>200MG/100ML (2MG/ML)</u>	<u>N050778 001</u> Sep 15, 1999
<u>AP</u>	+		<u>50MG/25ML (2MG/ML)</u>	<u>N050778 002</u> Sep 15, 1999

EPIRUBICIN HYDROCHLORIDE

<u>AP</u>		ACTAVIS TOTOWA	<u>50MG/25ML (2MG/ML)</u>	<u>A065445 002</u> Sep 18, 2008
<u>AP</u>			<u>200MG/100ML (2MG/ML)</u>	<u>A065445 003</u> Sep 18, 2008
<u>AP</u>		AKORN INC	<u>50MG/25ML (2MG/ML)</u>	<u>A090163 001</u> Jun 24, 2009
<u>AP</u>		CIPLA LTD	<u>50MG/25ML (2MG/ML)</u>	<u>A065361 001</u> Oct 22, 2007
<u>AP</u>			<u>200MG/100ML (2MG/ML)</u>	<u>A065361 002</u> Oct 22, 2007
<u>AP</u>		FRESENIUS KABI USA	<u>50MG/25ML (2MG/ML)</u>	<u>A065408 002</u> Oct 15, 2007
<u>AP</u>			<u>200MG/100ML (2MG/ML)</u>	<u>A065408 004</u> Oct 15, 2007
<u>AP</u>		HISUN PHARM HANGZHOU	<u>50MG/25ML (2MG/ML)</u>	<u>A090075 001</u> Mar 25, 2010
<u>AP</u>			<u>200MG/100ML (2MG/ML)</u>	<u>A090075 002</u> Mar 25, 2010
<u>AP</u>		IMPAX LABS INC	<u>50MG/25ML (2MG/ML)</u>	<u>A065331 001</u> Aug 09, 2007
<u>AP</u>			<u>200MG/100ML (2MG/ML)</u>	<u>A065331 002</u> Aug 09, 2007
<u>AP</u>		WEST-WARD PHARMS INT	<u>50MG/25ML (2MG/ML)</u>	<u>A065289 001</u> Jun 27, 2007
<u>AP</u>			<u>200MG/100ML (2MG/ML)</u>	<u>A065289 002</u> Jun 27, 2007
		FRESENIUS KABI USA	10MG/5ML (2MG/ML)	A065408 001 Oct 15, 2007
			150MG/75ML (2MG/ML)	A065408 003 Oct 15, 2007

EPLERENONE

TABLET; ORAL

EPLERENONE

<u>AB</u>		ACCORD HLTHCARE	<u>25MG</u>	<u>A206922 001</u> Jul 13, 2017
<u>AB</u>			<u>50MG</u>	<u>A206922 002</u> Jul 13, 2017
<u>AB</u>		BRECKENRIDGE	<u>25MG</u>	<u>A208283 001</u> Sep 14, 2018
<u>AB</u>			<u>50MG</u>	<u>A208283 002</u> Sep 14, 2018
<u>AB</u>		COREPHARMA	<u>25MG</u>	<u>A078482 001</u> Jul 30, 2008
<u>AB</u>			<u>50MG</u>	<u>A078482 002</u> Jul 30, 2008
<u>AB</u>		MYLAN	<u>25MG</u>	<u>A203896 001</u> Feb 02, 2017
<u>AB</u>			<u>50MG</u>	<u>A203896 002</u> Feb 02, 2017
<u>AB</u>		SANDOZ	<u>25MG</u>	<u>A078510 001</u> Aug 01, 2008
<u>AB</u>			<u>50MG</u>	<u>A078510 002</u> Aug 01, 2008
<u>INSPRA</u>				
<u>AB</u>	+	GD SEARLE LLC	<u>25MG</u>	<u>N021437 001</u> Sep 27, 2002
<u>AB</u>	+!		<u>50MG</u>	<u>N021437 002</u> Sep 27, 2002

EPOPROSTENOL SODIUM

INJECTABLE; INJECTION

EPOPROSTENOL SODIUM

<u>AP</u>		TEVA PHARMS USA	<u>EQ 0.5MG BASE/VIAL</u>	<u>A078396 001</u> Apr 23, 2008
<u>AP</u>			<u>EQ 1.5MG BASE/VIAL</u>	<u>A078396 002</u> Apr 23, 2008
<u>FLOLAN</u>				
<u>AP</u>	+!	GLAXOSMITHKLINE LLC	<u>EQ 0.5MG BASE/VIAL</u>	<u>N020444 001</u> Sep 20, 1995
<u>AP</u>	+!		<u>EQ 1.5MG BASE/VIAL</u>	<u>N020444 002</u> Sep 20, 1995
VELETRI				
	+	ACTELION PHARMS LTD	EQ 0.5MG BASE/VIAL	N022260 002 Jun 28, 2012
	+!		EQ 1.5MG BASE/VIAL	N022260 001 Jun 27, 2008

PRESCRIPTION DRUG PRODUCT LIST

3-163 (of 453)

EPROSARTAN MESYLATE

TABLET; ORAL

EPROSARTAN MESYLATE

<u>AB</u>	MYLAN PHARMS INC	<u>EQ 400MG BASE</u>	<u>A202012 001</u>	Nov 16, 2011
<u>AB</u>	!	<u>EQ 600MG BASE</u>	<u>A202012 002</u>	Nov 16, 2011

EPTIFIBATIDE

INJECTABLE; INJECTION

EPTIFIBATIDE

<u>AP</u>	ACCORD HLTHCARE	<u>2MG/ML</u>	<u>A205557 001</u>	Nov 06, 2017
<u>AP</u>		<u>75MG/100ML</u>	<u>A205557 002</u>	Nov 06, 2017
<u>AP</u>	AKORN	<u>2MG/ML</u>	<u>A204589 001</u>	Apr 18, 2017
<u>AP</u>		<u>75MG/100ML</u>	<u>A204589 002</u>	Apr 18, 2017
<u>AP</u>	AMNEAL PHARMS	<u>2MG/ML</u>	<u>A205581 001</u>	Dec 08, 2016
<u>AP</u>		<u>75MG/100ML</u>	<u>A205581 002</u>	Dec 08, 2016
<u>AP</u>	AUROBINDO PHARMA LTD	<u>2MG/ML</u>	<u>A206127 001</u>	Dec 08, 2015
<u>AP</u>		<u>75MG/100ML</u>	<u>A206127 002</u>	Dec 08, 2015
<u>AP</u>	BAXTER HLTHCARE CORP	<u>2MG/ML</u>	<u>A208554 001</u>	Nov 23, 2018
<u>AP</u>		<u>75MG/100ML</u>	<u>A208554 002</u>	Nov 23, 2018
<u>AP</u>	HAINAN POLY PHARM	<u>2MG/ML</u>	<u>A209864 001</u>	Jan 25, 2019
<u>AP</u>		<u>75MG/100ML</u>	<u>A209864 002</u>	Jan 25, 2019
<u>AP</u>	MYLAN LABS LTD	<u>2MG/ML</u>	<u>A203258 001</u>	Jul 20, 2018
<u>AP</u>		<u>75MG/100ML</u>	<u>A203258 002</u>	Jul 20, 2018
<u>AP</u>	SAGENT PHARMS INC	<u>2MG/ML</u>	<u>A204693 001</u>	Mar 07, 2018
<u>AP</u>		<u>75MG/100ML</u>	<u>A204693 002</u>	Mar 07, 2018
<u>AP</u>	TEVA PHARMS USA	<u>2MG/ML</u>	<u>A090854 001</u>	Jun 12, 2015
<u>INTEGRILIN</u>				
<u>AP</u>	+!	<u>2MG/ML</u>	<u>N020718 001</u>	May 18, 1998
<u>AP</u>	+	<u>75MG/100ML</u>	<u>N020718 002</u>	May 18, 1998

ERAVACYCLINE DIHYDROCHLORIDE

POWDER; INTRAVENOUS

XERAVA

+	TETRAPHASE PHARMS	EQ 50MG BASE/VIAL	N211109 001	Aug 27, 2018
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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

<u>AA</u>	+	US PHARM HOLDINGS	<u>50,000 IU</u>	<u>N003444 001</u>
<u>ERGOCALCIFEROL</u>				
<u>AA</u>		ORIT LABS LLC	<u>50,000 IU</u>	<u>A040833 001</u> May 20, 2009
<u>AA</u>		PURACAP PHARM LLC	<u>50,000 IU</u>	<u>A204276 001</u> Dec 07, 2018
<u>AA</u>		SIGMAPHARM LABS LLC	<u>50,000 IU</u>	<u>A091004 001</u> Jul 14, 2010
<u>AA</u>		STRIDES PHARMA	<u>50,000 IU</u>	<u>A090455 001</u> Aug 03, 2010
<u>AA</u>		SUN PHARM INDS INC	<u>50,000 IU</u>	<u>A040865 001</u> Dec 29, 2009
<u>VITAMIN D</u>				
<u>AA</u>		BIONPHARMA INC	<u>50,000 IU</u>	<u>A080704 001</u>

ERGOLOID MESYLATES

TABLET; ORAL

ERGOLOID MESYLATES

!	SUN PHARM INDUSTRIES	1MG	A081113 001	Oct 31, 1991
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ERGOTAMINE TARTRATE

TABLET; SUBLINGUAL

ERGOMAR

!	TERSERA THERAPS LLC	2MG	A087693 001	Feb 24, 1983
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ERIBULIN MESYLATE

SOLUTION; INTRAVENOUS

HALAVEN

+	EISAI INC	1MG/2ML (0.5MG/ML)	N201532 001	Nov 15, 2010
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PRESCRIPTION DRUG PRODUCT LIST

3-164 (of 453)

ERLOTINIB HYDROCHLORIDE

TABLET; ORAL

ERLOTINIB HYDROCHLORIDE

<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>	MYLAN	<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>	NATCO PHARMA LTD	<u>EQ 25MG BASE</u>	<u>A208488 001</u>	Nov 05, 2019
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A208488 002</u>	Nov 05, 2019
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A208488 003</u>	Nov 05, 2019
<u>AB</u>	SHILPA MEDICARE LTD	<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	<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>TARCEVA</u>				
<u>AB</u>	+	OSI PHARMS	<u>EQ 25MG BASE</u>	<u>N021743 001</u> Nov 18, 2004
<u>AB</u>	+		<u>EQ 100MG BASE</u>	<u>N021743 002</u> Nov 18, 2004
<u>AB</u>	+	!	<u>EQ 150MG BASE</u>	<u>N021743 003</u> Nov 18, 2004

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>	AUROBINDO PHARMA LTD	<u>EQ 1GM BASE/VIAL</u>	<u>A209133 001</u>	Jun 25, 2018
<u>AP</u>	SAVIOR LIFETEC CORP	<u>EQ 1GM BASE/VIAL</u>	<u>A207647 001</u>	Mar 19, 2019
<u>INVANZ</u>				
<u>AP</u>	+	MERCK SHARP DOHME	<u>EQ 1GM BASE/VIAL</u>	<u>N021337 001</u> Nov 21, 2001

ERTUGLIFLOZIN

TABLET; ORAL

STEGLATRO

+	MERCK SHARP DOHME	5MG	N209803 001	Dec 19, 2017
+	!	15MG	N209803 002	Dec 19, 2017

ERTUGLIFLOZIN; METFORMIN HYDROCHLORIDE

TABLET; ORAL

SEGLUROMET

+	MERCK SHARP DOHME	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

+	MERCK SHARP DOHME	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

<u>AB</u>	+	MAYNE PHARMA	<u>250MG</u>	<u>N050536 001</u>
<u>ERYTHROMYCIN</u>				
<u>AB</u>		ARBOR PHARMS LLC	<u>250MG</u>	<u>A062746 001</u> Dec 22, 1986

GEL; TOPICAL

ERYGEL

<u>AT</u>	+	MYLAN	<u>2%</u>	<u>N050617 001</u> Oct 21, 1987
<u>ERYTHROMYCIN</u>				
<u>AT</u>		FOUGERA PHARMS	<u>2%</u>	<u>A064184 001</u> Sep 30, 1997
<u>AT</u>		PERRIGO CO	<u>2%</u>	<u>A063211 001</u> Jan 29, 1993
<u>AT</u>		TELIGENT PHARMA INC	<u>2%</u>	<u>A208154 001</u> Jul 19, 2017

OINTMENT; OPHTHALMIC

ERYTHROMYCIN

<u>AT</u>		AKORN	<u>0.5%</u>	<u>A064030 001</u> Jul 18, 1996
<u>AT</u>		BAUSCH AND LOMB	<u>0.5%</u>	<u>A064067 001</u> Jul 29, 1994
<u>AT</u>	!	PERRIGO CO TENNESSEE	<u>0.5%</u>	<u>A062447 001</u> Sep 26, 1983

PRESCRIPTION DRUG PRODUCT LIST

ERYTHROMYCIN

SOLUTION; TOPICAL

ERYTHROMYCIN

AT	!	PERRIGO NEW YORK	2%	A063038	001	Jan 11, 1991
AT		TELLIGENT PHARMA INC	2%	A208100	001	Nov 20, 2017
AT		WOCKHARDT BIO AG	2%	A062825	001	Oct 23, 1987

SWAB; TOPICAL

ERYTHROMYCIN

AT		AKORN	2%	A090215	001	May 12, 2010
AT	!	PERRIGO CO	2%	A064126	001	Jul 03, 1996

TABLET; ORAL

ERYTHROMYCIN

AB		AMNEAL PHARMS CO	250MG	A209720	001	Mar 09, 2018
AB			500MG	A209720	002	Mar 09, 2018
AB		ARBOR PHARMS LLC	250MG	A061621	001	
AB	!		500MG	A061621	002	

TABLET, DELAYED RELEASE; ORAL

ERY-TAB

AB		ARBOR PHARMS LLC	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

ERYTHROMYCIN ETHYLSUCCINATE

GRANULE; ORAL

E.E.S.

AB	+	ARBOR PHARMS LLC	EQ 200MG BASE/5ML	N050207	001	
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ERYPED

AB	+	ARBOR PHARMS LLC	EQ 200MG BASE/5ML	N050207	003	Mar 30, 1987
AB	+	!	EQ 400MG BASE/5ML	N050207	002	

ERYTHROMYCIN ETHYLSUCCINATE

AB		AMNEAL PHARMS	EQ 200MG BASE/5ML	A211204	001	Nov 01, 2019
AB			EQ 400MG BASE/5ML	A211204	002	Nov 01, 2019
AB		ANI PHARMS INC	EQ 200MG BASE/5ML	A062055	001	
AB			EQ 200MG BASE/5ML	A062055	003	Nov 02, 2018
AB			EQ 400MG BASE/5ML	A062055	002	Nov 02, 2018
AB		PAR PHARM INC	EQ 200MG BASE/5ML	A211991	001	Oct 23, 2019
AB			EQ 400MG BASE/5ML	A211991	002	Oct 23, 2019

TABLET; ORAL

E.E.S. 400

BX	!	ARBOR PHARMS LLC	EQ 400MG BASE	A061905	002	Aug 12, 1982
BX	!	ARBOR PHARMS LLC	EQ 400MG BASE	A061904	001	

ERYTHROMYCIN LACTOBIONATE

INJECTABLE; INJECTION

ERYTHROCIN

AP		HOSPIRA	EQ 500MG BASE/VIAL	A062638	001	Oct 31, 1986
AP	+	!	EQ 500MG BASE/VIAL	N050609	001	Sep 24, 1986

ERYTHROMYCIN STEARATE

TABLET; ORAL

ERYTHROCIN STEARATE

!	ARBOR PHARMS LLC	EQ 250MG BASE	A060359	001	
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ESCITALOPRAM OXALATE

SOLUTION; ORAL

ESCITALOPRAM OXALATE

AA		AMNEAL PHARMS	EQ 5MG BASE/5ML	A202227	001	Mar 14, 2012
AA		ANTRIM PHARMS LLC	EQ 5MG BASE/5ML	A203967	001	May 26, 2015
AA		AUROBINDO PHARMA LTD	EQ 5MG BASE/5ML	A079062	001	Apr 02, 2012
AA		HETERO LABS LTD III	EQ 5MG BASE/5ML	A202221	001	Jun 12, 2012
AA		LANNETT CO INC	EQ 5MG BASE/5ML	A090477	001	Jun 12, 2013
AA		MACLEODS PHARMS LTD	EQ 5MG BASE/5ML	A202754	001	Mar 31, 2016
AA		TARO	EQ 5MG BASE/5ML	A079121	001	May 03, 2012

LEXAPRO

AA	+	ALLERGAN	EQ 5MG BASE/5ML	N021365	001	Nov 27, 2002
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TABLET; ORAL

ESCITALOPRAM OXALATE

AB		ACCORD HLTHCARE	EQ 5MG BASE	A202389	001	Sep 11, 2012
AB			EQ 10MG BASE	A202389	002	Sep 11, 2012

PRESCRIPTION DRUG PRODUCT LIST

ESCITALOPRAM OXALATE

TABLET; ORAL

ESCITALOPRAM OXALATE

<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>	CADILA	<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
<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>	HIKMA PHARMS	<u>EQ 5MG BASE</u>	<u>A078766 001</u>	Sep 11, 2012
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A078766 002</u>	Sep 11, 2012
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A078766 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 GENERICS	<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>	LUPIN LTD	<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>	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>	PHARM ASSOC	<u>EQ 5MG BASE</u>	<u>A077512 001</u>	Sep 12, 2012
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A077512 002</u>	Sep 12, 2012
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A077512 003</u>	Sep 12, 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
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A078032 003</u>	Aug 28, 2015
<u>AB</u>	TEVA PHARMS USA	<u>EQ 5MG BASE</u>	<u>A076765 001</u>	Mar 14, 2012
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A076765 002</u>	Mar 14, 2012
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A076765 003</u>	Mar 14, 2012
<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>LEXAPRO</u>			
<u>AB</u>	+ ALLERGAN	<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

+	SUNOVION PHARMS INC	200MG	N022416 001	Nov 08, 2013
+		400MG	N022416 002	Nov 08, 2013
+		600MG	N022416 003	Nov 08, 2013
+		800MG	N022416 004	Nov 08, 2013

ESMOLOL HYDROCHLORIDE

INJECTABLE; INJECTION

BREVIBLOC

<u>AP</u>	+! BAXTER HLTHCARE	<u>10MG/ML</u>	<u>N019386 006</u>	Feb 25, 2003
	<u>ESMOLOL HYDROCHLORIDE</u>			
<u>AP</u>	AUROBINDO PHARMA LTD	<u>10MG/ML</u>	<u>A205520 001</u>	Jul 23, 2015
<u>AP</u>	FRESENIUS KABI USA	<u>10MG/ML</u>	<u>A076573 001</u>	May 02, 2005
<u>AP</u>	GLAND PHARMA LTD	<u>10MG/ML</u>	<u>A208538 001</u>	Aug 14, 2019
<u>AP</u>	MYLAN INSTITUTIONAL	<u>10MG/ML</u>	<u>A076474 001</u>	May 02, 2005
<u>AP</u>	MYLAN LABS LTD	<u>10MG/ML</u>	<u>A206608 001</u>	Jun 08, 2018
<u>AP</u>		<u>20MG/ML</u>	<u>A206608 002</u>	Jun 08, 2018

PRESCRIPTION DRUG PRODUCT LIST

ESMOLOL HYDROCHLORIDE

INJECTABLE; INJECTION

ESMOLOL HYDROCHLORIDE

AP	SAGENT PHARMS INC	<u>10MG/ML</u>	<u>A207107</u>	<u>001</u>	Jun 08, 2018
AP		<u>20MG/ML</u>	<u>A207107</u>	<u>002</u>	Jun 08, 2018
AP	WEST-WARD PHARMS INT	<u>10MG/ML</u>	<u>A076323</u>	<u>001</u>	Aug 10, 2004
	BREVIBLOC DOUBLE STRENGTH IN PLASTIC CONTAINER				
	+! BAXTER HLTHCARE	2GM/100ML	N019386	005	Jan 27, 2003
	BREVIBLOC IN PLASTIC CONTAINER				
	+! BAXTER HLTHCARE	1GM/100ML	N019386	004	Feb 16, 2001
	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

ESOMEPRAZOLE MAGNESIUM

CAPSULE, DELAYED REL PELLETS; ORAL

ESOMEPRAZOLE MAGNESIUM

AB	ALKEM LABS LTD	<u>EQ 20MG BASE</u>	<u>A208333</u>	<u>001</u>	Oct 20, 2017
AB		<u>EQ 40MG BASE</u>	<u>A208333</u>	<u>002</u>	Oct 20, 2017
AB	AUROBINDO PHARMA LTD	<u>EQ 20MG BASE</u>	<u>A205606</u>	<u>001</u>	Apr 21, 2016
AB		<u>EQ 40MG BASE</u>	<u>A205606</u>	<u>002</u>	Apr 21, 2016
AB	DR REDDYS LABS LTD	<u>EQ 20MG BASE</u>	<u>A078279</u>	<u>001</u>	Sep 25, 2015
AB		<u>EQ 40MG BASE</u>	<u>A078279</u>	<u>002</u>	Sep 25, 2015
AB	GLENMARK PHARMS	<u>EQ 20MG BASE</u>	<u>A209495</u>	<u>001</u>	May 10, 2019
AB		<u>EQ 40MG BASE</u>	<u>A209495</u>	<u>002</u>	May 10, 2019
AB	HETERO LABS LTD III	<u>EQ 20MG BASE</u>	<u>A202784</u>	<u>001</u>	Sep 21, 2015
AB		<u>EQ 40MG BASE</u>	<u>A202784</u>	<u>002</u>	Sep 21, 2015
AB	IVAX SUB TEVA PHARMS	<u>EQ 20MG BASE</u>	<u>A078003</u>	<u>001</u>	Jan 26, 2015
AB		<u>EQ 40MG BASE</u>	<u>A078003</u>	<u>002</u>	Jan 26, 2015
AB	LANNETT CO INC	<u>EQ 20MG BASE</u>	<u>A205563</u>	<u>001</u>	Sep 01, 2017
AB		<u>EQ 40MG BASE</u>	<u>A205563</u>	<u>002</u>	Sep 01, 2017
AB	MYLAN	<u>EQ 20MG BASE</u>	<u>A078936</u>	<u>001</u>	Aug 02, 2015
AB		<u>EQ 40MG BASE</u>	<u>A078936</u>	<u>002</u>	Aug 03, 2015
AB	SUN PHARM	<u>EQ 20MG BASE</u>	<u>A209735</u>	<u>001</u>	Apr 30, 2018
AB		<u>EQ 40MG BASE</u>	<u>A209735</u>	<u>002</u>	Apr 30, 2018
AB	TORRENT	<u>EQ 20MG BASE</u>	<u>A203636</u>	<u>001</u>	Oct 19, 2015
AB		<u>EQ 40MG BASE</u>	<u>A203636</u>	<u>002</u>	Oct 19, 2015
AB	ZYDUS PHARMS	<u>EQ 20MG BASE</u>	<u>A206296</u>	<u>001</u>	May 22, 2019
AB		<u>EQ 40MG BASE</u>	<u>A206296</u>	<u>002</u>	May 22, 2019

NEXIUM

AB	+ ASTRAZENECA PHARMS	<u>EQ 20MG BASE</u>	<u>N021153</u>	<u>001</u>	Feb 20, 2001
AB	+!	<u>EQ 40MG BASE</u>	<u>N021153</u>	<u>002</u>	Feb 20, 2001

FOR SUSPENSION, DELAYED RELEASE; ORAL

NEXIUM

	+ ASTRAZENECA PHARMS	EQ 2.5MG BASE/PACKET	N021957	003	Dec 15, 2011
		EQ 5MG BASE/PACKET	N021957	004	Dec 15, 2011
		EQ 10MG BASE/PACKET	N022101	001	Feb 27, 2008
		EQ 20MG BASE/PACKET	N021957	001	Oct 20, 2006
	+!	EQ 40MG BASE/PACKET	N021957	002	Oct 20, 2006

ESOMEPRAZOLE MAGNESIUM; NAPROXEN

TABLET, DELAYED RELEASE; ORAL

NAPROXEN AND ESOMEPRAZOLE MAGNESIUM

AB	DR REDDYS LABS LTD	<u>EQ 20MG BASE; 375MG</u>	<u>A204206</u>	<u>001</u>	Feb 18, 2020
AB		<u>EQ 20MG BASE; 500MG</u>	<u>A204206</u>	<u>002</u>	Feb 18, 2020
	<u>VIMOVO</u>				
AB	+ HORIZON	<u>EQ 20MG BASE; 375MG</u>	<u>N022511</u>	<u>002</u>	Apr 30, 2010
AB	+!	<u>EQ 20MG BASE; 500MG</u>	<u>N022511</u>	<u>001</u>	Apr 30, 2010

ESOMEPRAZOLE SODIUM

INJECTABLE; INTRAVENOUS

ESOMEPRAZOLE SODIUM

AP	ACCORD HLTHCARE	<u>EQ 40MG BASE/VIAL</u>	<u>A205379</u>	<u>001</u>	Sep 25, 2015
AP	! AUROBINDO PHARMA LTD	<u>EQ 40MG BASE/VIAL</u>	<u>A204657</u>	<u>002</u>	Aug 10, 2016
AP	DEVA HOLDING AS	<u>EQ 40MG BASE/VIAL</u>	<u>A207181</u>	<u>001</u>	Mar 06, 2017
AP	MYLAN LABS LTD	<u>EQ 40MG BASE/VIAL</u>	<u>A202686</u>	<u>002</u>	May 17, 2017
AP	SUN PHARMA GLOBAL	<u>EQ 40MG BASE/VIAL</u>	<u>A200882</u>	<u>002</u>	Mar 18, 2013

PRESCRIPTION DRUG PRODUCT LIST

ESOMEPRAZOLE SODIUM

INJECTABLE; INTRAVENOUS

NEXIUM IV

<u>AP</u>	<u>+!</u>	ASTRAZENECA PHARMS	<u>EO 40MG BASE/VIAL</u>	<u>N021689 002</u>	Mar 31, 2005
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ESTAZOLAM

TABLET; ORAL

ESTAZOLAM

<u>AB</u>		MAYNE PHARMA	<u>1MG</u>	<u>A074921 001</u>	Jul 10, 1997
<u>AB</u>	<u>!</u>		<u>2MG</u>	<u>A074921 002</u>	Jul 10, 1997
<u>AB</u>		PAR PHARM	<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>	<u>!</u>	ALLERGAN	<u>0.01%</u>	<u>A086069 001</u>	Jan 31, 1984
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ESTRADIOL

<u>AB</u>		ALVOGEN PINE BROOK	<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>		PERRIGO UK FINCO	<u>0.01%</u>	<u>A210194 001</u>	Jan 22, 2018
<u>AB</u>		TEVA PHARMS USA	<u>0.01%</u>	<u>A210488 001</u>	Mar 30, 2018

FILM, EXTENDED RELEASE; TRANSDERMAL

CLIMARA

<u>AB</u>	<u>+</u>	BAYER HLTHCARE	<u>0.06MG/24HR</u>	<u>N020375 006</u>	May 27, 2003
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ESTRADIOL

<u>AB</u>		MYLAN TECHNOLOGIES	<u>0.06MG/24HR</u>	<u>A075182 005</u>	Jul 20, 2006
<u>AB1</u>		AMNEAL PHARMS LLC	<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

VIVELLE-DOT

<u>AB1</u>	<u>+</u>	NOVARTIS	<u>0.025MG/24HR</u>	<u>N020538 009</u>	May 03, 2002
<u>AB1</u>	<u>+</u>		<u>0.0375MG/24HR</u>	<u>N020538 005</u>	Jan 08, 1999
<u>AB1</u>	<u>+</u>		<u>0.05MG/24HR</u>	<u>N020538 006</u>	Jan 08, 1999
<u>AB1</u>	<u>+</u>		<u>0.075MG/24HR</u>	<u>N020538 007</u>	Jan 08, 1999
<u>AB1</u>	<u>+!</u>		<u>0.1MG/24HR</u>	<u>N020538 008</u>	Jan 08, 1999

CLIMARA

<u>AB2</u>	<u>+</u>	BAYER HLTHCARE	<u>0.025MG/24HR</u>	<u>N020375 004</u>	Mar 05, 1999
<u>AB2</u>	<u>+</u>		<u>0.0375MG/24HR</u>	<u>N020375 005</u>	May 27, 2003
<u>AB2</u>	<u>+</u>		<u>0.05MG/24HR</u>	<u>N020375 001</u>	Dec 22, 1994
<u>AB2</u>	<u>+</u>		<u>0.075MG/24HR</u>	<u>N020375 003</u>	Mar 23, 1998
<u>AB2</u>	<u>+!</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>0.075MG/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>AB3</u>			<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>	<u>+</u>	NOVEN	<u>0.025MG/24HR</u>	<u>N203752 005</u>	Sep 23, 2014
<u>AB3</u>	<u>+</u>		<u>0.0375MG/24HR</u>	<u>N203752 001</u>	Oct 29, 2012
<u>AB3</u>	<u>+</u>		<u>0.05MG/24HR</u>	<u>N203752 003</u>	Oct 29, 2012
<u>AB3</u>	<u>+</u>		<u>0.075MG/24HR</u>	<u>N203752 002</u>	Oct 29, 2012
<u>AB3</u>	<u>+!</u>		<u>0.1MG/24HR</u>	<u>N203752 004</u>	Oct 29, 2012

ALORA

BX		ALLERGAN	0.025MG/24HR	N020655 004	Apr 05, 2002
BX			0.05MG/24HR	N020655 001	Dec 20, 1996
BX			0.075MG/24HR	N020655 002	Dec 20, 1996
BX			0.1MG/24HR	N020655 003	Dec 20, 1996

PRESCRIPTION DRUG PRODUCT LIST

ESTRADIOL

FILM, EXTENDED RELEASE;TRANSDERMAL

MENOSTAR

+! BAYER HLTHCARE 0.014MG/24HR N021674 001 Jun 08, 2004

GEL;TRANSDERMAL

DIVIGEL

+! VERTICAL PHARMS LLC 0.1% N022038 001 Jun 04, 2007

GEL, METERED;TRANSDERMAL

ELESTRIN

+! MYLAN SPECIALITY LP 0.06% (0.87GM/ACTIVATION) N021813 001 Dec 15, 2006

ESTROGEL

+! ASCEND THERAPS US 0.06% (1.25GM/ACTIVATION) N021166 002 Feb 09, 2004

INSERT;VAGINAL

IMVEXXY

+ THERAPEUTICSMD INC 0.004MG N208564 001 May 29, 2018

+! 0.01MG N208564 002 May 29, 2018

INSERT, EXTENDED RELEASE;VAGINAL

ESTRING

+! PHARMACIA AND 0.0075MG/24HR N020472 001 Apr 26, 1996

UPJOHN

SPRAY;TRANSDERMAL

EVAMIST

+! PERRIGO PHARMA INTL 1.53MG/SPRAY N022014 001 Jul 27, 2007

TABLET;ORAL

ESTRADIOL

AB	BARR LABS INC	0.5MG	A040197 001	Oct 22, 1997
AB		1MG	A040197 002	Oct 22, 1997
AB	!	2MG	A040197 003	Oct 22, 1997
AB	EPIC PHARMA INC	0.5MG	A040275 001	Dec 29, 1998
AB		1MG	A040275 002	Dec 29, 1998
AB		2MG	A040275 003	Dec 29, 1998
AB	MAYNE PHARMA	0.5MG	A040114 003	Mar 14, 1996
AB		1MG	A040114 001	Mar 14, 1996
AB		2MG	A040114 002	Mar 14, 1996
AB	MYLAN	0.5MG	A040326 001	Apr 21, 1999
AB		1MG	A040326 002	Apr 21, 1999
AB		2MG	A040326 003	Apr 21, 1999

TABLET;VAGINAL

ESTRADIOL

AB	AMNEAL PHARMS	10MCG	A205256 001	May 29, 2015
AB	GLENMARK PHARMS LTD	10MCG	A210264 001	Sep 14, 2018
AB	TEVA PHARMS USA	10MCG	A206388 001	Jul 21, 2017

VAGIFEM

AB	+! NOVO NORDISK INC	10MCG	N020908 002	Nov 25, 2009
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ESTRADIOL ACETATE

INSERT, EXTENDED RELEASE;VAGINAL

FEMRING

+ MILLICENT 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

! PHARMACIA AND 5MG/ML A085470 003

UPJOHN

ESTRADIOL VALERATE

INJECTABLE;INJECTION

DELESTROGEN

AO	+! PAR STERILE PRODUCTS	20MG/ML	N009402 004	
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AO	+!	40MG/ML	N009402 003	
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ESTRADIOL VALERATE

AO	AM REGENT	20MG/ML	A090920 001	Jan 19, 2010
AO		40MG/ML	A090920 002	Jan 19, 2010

DELESTROGEN

+! PAR STERILE 10MG/ML N009402 002

PRODUCTS

PRESCRIPTION DRUG PRODUCT LISTESTRADIOL; 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**AB** + AMNEAL PHARMS LLC **0.5MG;0.1MG** **N020907 002** Dec 28, 2006**AB** +! **1MG;0.5MG** **N020907 001** Nov 18, 1998AMABELZ**AB** LUPIN LTD **0.5MG;0.1MG** **A203339 001** Jun 20, 2016**AB** **1MG;0.5MG** **A203339 002** Jun 20, 2016ESTRADIOL AND NORETHINDRONE ACETATE**AB** BARR **1MG;0.5MG** **A079193 001** May 11, 2010**AB** BRECKENRIDGE PHARM **0.5MG;0.1MG** **A078324 002** Jun 09, 2011**AB** **1MG;0.5MG** **A078324 001** Apr 17, 2008**AB** MYLAN LABS LTD **0.5MG;0.1MG** **A207261 001** Feb 10, 2017**AB** **1MG;0.5MG** **A207261 002** Feb 10, 2017**AB** NOVAST LABS **0.5MG;0.1MG** **A210612 001** Apr 03, 2019**AB** **1MG;0.5MG** **A210612 002** Apr 03, 2019ESTRADIOL; NORGESTIMATE

TABLET;ORAL

ESTRADIOL AND NORGESTIMATE

! BARR 1MG,1MG;N/A,0.09MG A076812 001 Apr 29, 2005

ESTRADIOL; PROGESTERONE

CAPSULE;ORAL

BIJUVA

+! THERAPEUTICSMD INC 1MG;100MG N210132 001 Oct 28, 2018

ESTRAMUSTINE PHOSPHATE SODIUM

CAPSULE;ORAL

EMCYT

+! PHARMACIA AND UPJOHN EQ 140MG PHOSPHATE N018045 001

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

ESTROGENS, ESTERIFIED

TABLET;ORAL

MENEST

MONARCH PHARMS 0.3MG A084951 001

0.625MG A084948 001

1.25MG A084950 001

! 2.5MG A084949 001

PRESCRIPTION DRUG PRODUCT LIST

ESTROPIPATE

TABLET; ORAL

OGEN 5

PHARMACIA AND
UPJOHN

6MG

A083220 004

ESZOPICLONE

TABLET; ORAL

ESZOPICLONE

AB	AUROBINDO PHARMA LTD	1MG	A208451 001	Sep 15, 2016
AB		2MG	A208451 002	Sep 15, 2016
AB		3MG	A208451 003	Sep 15, 2016
AB	DR REDDYS LABS LTD	1MG	A091024 001	Apr 15, 2014
AB		2MG	A091024 002	Apr 15, 2014
AB		3MG	A091024 003	Apr 15, 2014
AB	GLENMARK GENERICS	1MG	A091166 001	Apr 15, 2014
AB		2MG	A091166 002	Apr 15, 2014
AB		3MG	A091166 003	Apr 15, 2014
AB	LUPIN LTD	1MG	A091124 001	Sep 13, 2011
AB		2MG	A091124 002	Sep 13, 2011
AB		3MG	A091124 003	Sep 13, 2011
AB	MACLEODS PHARMS LTD	1MG	A202929 001	Jan 30, 2015
AB		2MG	A202929 002	Jan 30, 2015
AB		3MG	A202929 003	Jan 30, 2015
AB	MYLAN PHARMS INC	1MG	A091151 001	Mar 26, 2013
AB		2MG	A091151 002	Mar 26, 2013
AB		3MG	A091151 003	Mar 26, 2013
AB	ORCHID HLTHCARE	1MG	A091113 001	Jun 10, 2014
AB		2MG	A091113 002	Jun 10, 2014
AB		3MG	A091113 003	Jun 10, 2014
AB	SUN PHARM	1MG	A091103 001	Apr 03, 2013
AB		2MG	A091103 002	Apr 03, 2013
AB		3MG	A091103 003	Apr 03, 2013
AB	TEVA	1MG	A091169 001	May 23, 2011
AB		2MG	A091169 002	May 23, 2011
AB		3MG	A091169 003	May 23, 2011
	LUNESTA			
AB	+ SUNOVION PHARMS INC	1MG	N021476 001	Dec 15, 2004
AB	+	2MG	N021476 002	Dec 15, 2004
AB	+!	3MG	N021476 003	Dec 15, 2004

ETELCACTIDE

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

EDECRIIN

AP	+!	BAUSCH	EQ 50MG BASE/VIAL	N016093 001	
		ETHACRYNATE SODIUM			
AP		MYLAN INSTITUTIONAL	EQ 50MG BASE/VIAL	A204634 001	Aug 23, 2016
AP		PAR STERILE PRODUCTS	EQ 50MG BASE/VIAL	A205473 001	Jul 29, 2015
AP		ZYDUS PHARMS	EQ 50MG BASE/VIAL	A207758 001	Nov 17, 2017

ETHACRYNIC ACID

TABLET; ORAL

EDECRIIN

AB	+!	BAUSCH	25MG	N016092 001	
		ETHACRYNIC ACID			
AB		AGNITIO	25MG	A211809 001	Jul 12, 2019
AB		ALVOGEN	25MG	A205709 001	Jul 24, 2018
AB		AMNEAL PHARMS CO	25MG	A208805 001	May 08, 2018
AB		EDENBRIDGE PHARMS	25MG	A205609 001	Jun 30, 2016
AB		HIKMA	25MG	A207262 001	Feb 23, 2017

PRESCRIPTION DRUG PRODUCT LIST

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ETHACRYNIC ACID

TABLET; ORAL

ETHACRYNIC ACID

AB	LUPIN LTD	25MG	A211719 001	Sep 06, 2019
AB	PAR PHARM INC	25MG	A208501 001	Jul 21, 2017
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	AKORN	100MG	A075095 001	Nov 30, 1999
AB		400MG	A075095 002	Nov 30, 1999
AB	BARR	400MG	A076057 001	Nov 26, 2001
AB	LUPIN	100MG	A078939 001	Jun 17, 2009
AB		400MG	A078939 002	Jun 17, 2009

MYAMBUTOL

AB	+ STI PHARMA LLC	100MG	N016320 001	
AB	+!	400MG	N016320 003	

ETHANOLAMINE OLEATE

INJECTABLE; INJECTION

ETHAMOLIN

+!	QOL MEDCL	50MG/ML	N019357 001	Dec 22, 1988
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ETHINYL ESTRADIOL; ETHYNODIOL DIACETATE

TABLET; ORAL-28

ETHYNODIOL DIACETATE AND ETHINYL ESTRADIOL

AB	MYLAN LABS LTD	0.035MG;1MG	A204703 001	Jul 28, 2016
AB		0.05MG;1MG	A204704 001	Feb 09, 2016
	KELNOR			
AB	BARR	0.035MG;1MG	A076785 001	May 23, 2005
	LO-MALMOREDE			
AB	NOVAST LABS	0.035MG;1MG	A209548 001	Feb 11, 2019
	MALMOREDE			
AB	NOVAST LABS	0.05MG;1MG	A209547 001	Jul 25, 2018
	ZOVIA 1/35E-28			
AB	MAYNE PHARMA	0.035MG;1MG	A072721 001	Dec 30, 1991
	ZOVIA 1/50E-28			
AB	! WATSON LABS	0.05MG;1MG	A072723 001	Dec 30, 1991

ETHINYL ESTRADIOL; ETONOGESTREL

RING; VAGINAL

ELURYNG

AB	AMNEAL PHARMS LLC	0.015MG/24HR;0.12MG/24HR	A210830 001	Dec 11, 2019
	NUVARING			
AB	+! ORGANON SUB MERCK	0.015MG/24HR;0.12MG/24HR	N021187 001	Oct 03, 2001

ETHINYL ESTRADIOL; LEVONORGESTREL

SYSTEM; TRANSDERMAL

TWIRLA

+!	AGILE	0.03MG/24HR;0.12MG/24HR	N204017 001	Feb 14, 2020
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TABLET; ORAL

ASHLYNA

AB	GLENMARK GENERICS	0.03MG,0.01MG;0.15MG,N/A	A203163 001	Feb 23, 2015
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DAYSEE

AB	LUPIN LTD	0.03MG,0.01MG;0.15MG,N/A	A091467 001	Apr 10, 2013
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FAYOSIM

AB	LUPIN LTD	0.02MG,0.15MG;0.025MG,0.15MG;0.03MG,0.15MG;0.01MG,N/A	A205943 001	Mar 29, 2016
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ICLEVIA

AB	AUROBINDO PHARMA LTD	0.03MG;0.15MG	A206850 001	Jun 29, 2018
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INTROVALE

AB	XIROMED	0.03MG;0.15MG	A079064 001	Sep 27, 2010
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JAIMIESS

AB	XIROMED	0.03MG,0.01MG;0.15MG,N/A	A203770 001	Dec 27, 2017
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LEVONORGESTREL AND ETHINYL ESTRADIOL

AB	AMNEAL PHARMS	0.03MG;0.15MG	A203871 001	Nov 13, 2015
AB		0.03MG,0.01MG;0.15MG,N/A	A203872 001	Dec 22, 2015
AB	GLENMARK GENERICS	0.02MG;0.09MG	A202791 001	Apr 09, 2015
AB	GLENMARK PHARMS LTD	0.03MG;0.15MG	A203164 001	Jun 12, 2015
AB	LUPIN LTD	0.03MG;0.15MG	A091440 001	Oct 23, 2012
AB	MYLAN LABS LTD	0.03MG;0.15MG	A200490 001	Apr 21, 2015
AB	! WATSON LABS	0.02MG;0.09MG	A079218 001	Jun 06, 2011

PRESCRIPTION DRUG PRODUCT LIST

ETHINYL ESTRADIOL; LEVONORGESTREL

TABLET; ORAL

LEVONORGESTREL AND ETHINYL ESTRADIOL AND ETHINYL ESTRADIOL

<u>AB</u>	LUPIN LTD	<u>0.02MG, 0.1MG; 0.01MG, N/A</u>	<u>A091674 001</u>	Oct 26, 2011
<u>AB</u>	MAYNE PHARMA	<u>0.02MG, 0.1MG; 0.01MG, N/A</u>	<u>A200407 001</u>	Oct 25, 2011
<u>AB</u>		<u>0.03MG, 0.01MG; 0.15MG, N/A</u>	<u>A078834 001</u>	May 31, 2011
<u>AB</u>	MYLAN LABS LTD	<u>0.02MG, 0.15MG;</u> <u>0.025MG, 0.15MG; 0.03MG, 0.15MG;</u>	<u>A206053 001</u>	Oct 02, 2017
<u>AB</u>		<u>0.02MG, 0.1MG; 0.01MG, N/A</u>	<u>A200493 001</u>	Jun 17, 2015
<u>AB</u>		<u>0.03MG, 0.01MG; 0.15MG, N/A</u>	<u>A200492 001</u>	May 27, 2015
<u>AB</u>	XIROMED	<u>0.02MG, 0.1MG; 0.01MG, N/A</u>	<u>A205131 001</u>	Dec 14, 2017
	<u>LO SIMPESSE</u>			
<u>AB</u>	AUROBINDO PHARMA LTD	<u>0.02MG, 0.1MG; 0.01MG, N/A</u>	<u>A206852 001</u>	Apr 28, 2017
	<u>LOSEASONIQUE</u>			
<u>AB</u>	TEVA BRANDED PHARM	<u>0.02MG, 0.1MG; 0.01MG, N/A</u>	<u>N022262 001</u>	Oct 24, 2008
	<u>QUARTETTE</u>			
<u>AB</u>	+! TEVA BRANDED PHARM	<u>0.02MG, 0.15MG; 0.025MG, 0.15MG; 0.03MG, 0.15MG; 0.01MG, N/A</u>	<u>N204061 001</u>	Mar 28, 2013
	<u>QUASENSE</u>			
<u>AB</u>	WATSON LABS	<u>0.03MG; 0.15MG</u>	<u>A077101 001</u>	Sep 06, 2006
	<u>SEASONALE</u>			
<u>AB</u>	+! TEVA BRANDED PHARM	<u>0.03MG; 0.15MG</u>	<u>N021544 001</u>	Sep 05, 2003
	<u>SEASONIQUE</u>			
<u>AB</u>	+! TEVA BRANDED PHARM	<u>0.03MG, 0.01MG; 0.15MG, N/A</u>	<u>N021840 001</u>	May 25, 2006
	<u>SETLAKIN</u>			
<u>AB</u>	NOVAST LABS	<u>0.03MG; 0.15MG</u>	<u>A090716 001</u>	Sep 15, 2014
	<u>SIMPESSE</u>			
<u>AB</u>	AUROBINDO PHARMA LTD	<u>0.03MG, 0.01MG; 0.15MG, N/A</u>	<u>A206851 001</u>	Apr 07, 2017
	<u>SYLEVIA</u>			
<u>AB</u>	SUN PHARM	<u>0.03MG; 0.15MG</u>	<u>A202988 001</u>	Feb 06, 2019
	BALCOLTRA			
	+! AVION PHARMS	0.02MG; 0.1MG	N208612 001	Jan 09, 2018
	TABLET; ORAL-28			
	<u>ALTAVERA</u>			
<u>AB</u>	XIROMED	<u>0.03MG; 0.15MG</u>	<u>A079102 001</u>	Aug 03, 2010
	<u>AYUNA</u>			
<u>AB</u>	AUROBINDO PHARMA LTD	<u>0.03MG; 0.15MG</u>	<u>A206866 001</u>	Sep 23, 2016
	<u>ELIFEMME</u>			
<u>AB</u>	XIROMED	<u>0.03MG, 0.04MG, 0.03MG; 0.05MG, 0.075MG, 0.125MG</u>	<u>A202507 001</u>	Dec 04, 2015
	<u>ENPRESSE-28</u>			
<u>AB</u>	DURAMED PHARMS BARR	<u>0.03MG, 0.04MG, 0.03MG; 0.05MG, 0.075MG, 0.125MG</u>	<u>A075809 002</u>	Jul 16, 2001
	<u>KURVELO</u>			
<u>AB</u>	LUPIN LTD	<u>0.03MG; 0.15MG</u>	<u>A091408 001</u>	Oct 17, 2012
	<u>LEVONEST</u>			
<u>AB</u>	NOVAST LABS LTD	<u>0.03MG, 0.04MG, 0.03MG; 0.05MG, 0.075MG, 0.125MG</u>	<u>A090719 001</u>	Dec 29, 2010
	<u>LEVONORGESTREL AND ETHINYL ESTRADIOL</u>			
<u>AB</u>	AMNEAL PHARMS	<u>0.03MG; 0.15MG</u>	<u>A201095 001</u>	Dec 08, 2014
<u>AB</u>	LUPIN LTD	<u>0.03MG, 0.04MG, 0.03MG; 0.05MG, 0.075MG, 0.125MG</u>	<u>A200248 001</u>	Nov 19, 2015
<u>AB</u>	MYLAN LABS LTD	<u>0.03MG; 0.15MG</u>	<u>A091663 001</u>	Dec 21, 2012
	<u>LEVORA 0.15/30-28</u>			
<u>AB</u>	! MAYNE PHARMA	<u>0.03MG; 0.15MG</u>	<u>A073594 001</u>	Dec 13, 1993
	<u>MARLISSA</u>			
<u>AB</u>	GLENMARK GENERICS	<u>0.03MG; 0.15MG</u>	<u>A091452 001</u>	Feb 29, 2012
	<u>MYZILRA</u>			
<u>AB</u>	VINTAGE PHARMS LLC	<u>0.03MG, 0.04MG, 0.03MG; 0.05MG, 0.075MG, 0.125MG</u>	<u>A077502 001</u>	Nov 23, 2011
	<u>PORTIA-28</u>			
<u>AB</u>	BARR	<u>0.03MG; 0.15MG</u>	<u>A075866 002</u>	May 23, 2002
	<u>TRIVORA-28</u>			
<u>AB</u>	! MAYNE PHARMA	<u>0.03MG, 0.04MG, 0.03MG; 0.05MG, 0.075MG, 0.125MG</u>	<u>A074538 002</u>	Dec 18, 1997
	<u>AFIRMELLE</u>			
<u>AB1</u>	AUROBINDO PHARMA LTD	<u>0.02MG; 0.1MG</u>	<u>A206886 001</u>	Nov 14, 2016
	<u>AVIANE-28</u>			
<u>AB1</u>	DURAMED PHARMS BARR	<u>0.02MG; 0.1MG</u>	<u>A075796 001</u>	Apr 30, 2001

PRESCRIPTION DRUG PRODUCT LIST

ETHINYL ESTRADIOL; LEVONORGESTREL

TABLET; ORAL-28

CERINTA

AB1	SUN PHARM	<u>0.02MG;0.1MG</u>	<u>A202817</u>	<u>001</u>	Jan 07, 2019
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FALMINA

AB1	NOVAST LABS LTD	<u>0.02MG;0.1MG</u>	<u>A090721</u>	<u>001</u>	Mar 28, 2012
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LEVONORGESTREL AND ETHINYL ESTRADIOL

AB1	AMNEAL PHARMS	<u>0.02MG;0.1MG</u>	<u>A201108</u>	<u>001</u>	Feb 05, 2014
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AB1	LUPIN LTD	<u>0.02MG;0.1MG</u>	<u>A091425</u>	<u>001</u>	Jan 18, 2013
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AB1	! MAYNE PHARMA	<u>0.02MG;0.1MG</u>	<u>A076625</u>	<u>001</u>	Nov 18, 2004
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AB1	MYLAN LABS LTD	<u>0.02MG;0.1MG</u>	<u>A200245</u>	<u>001</u>	Oct 09, 2013
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ORSYTHIA

AB1	VINTAGE PHARMS LLC	<u>0.02MG;0.1MG</u>	<u>A077099</u>	<u>001</u>	May 11, 2011
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VIENVA

AB1	XIROMED	<u>0.02MG;0.1MG</u>	<u>A201088</u>	<u>001</u>	May 21, 2015
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LESSINA-28

AB2	BARR	<u>0.02MG;0.1MG</u>	<u>A075803</u>	<u>002</u>	Mar 20, 2002
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LEVONORGESTREL AND ETHINYL ESTRADIOL

AB2	! MAYNE PHARMA	<u>0.02MG;0.1MG</u>	<u>A077681</u>	<u>001</u>	May 31, 2006
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ETHINYL ESTRADIOL; NORELGESTROMIN

FILM, EXTENDED RELEASE; TRANSDERMAL

XULANE

!	MYLAN TECHNOLOGIES	<u>0.035MG/24HR;0.15MG/24HR</u>	A200910	001	Apr 16, 2014
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ETHINYL ESTRADIOL; NORETHINDRONE

TABLET; ORAL-21

NORINYL 1+35 21-DAY

AB	ALLERGAN	<u>0.035MG;1MG</u>	<u>N017565</u>	<u>001</u>	
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NORTREL 1/35-21

AB	BARR	<u>0.035MG;1MG</u>	<u>A072693</u>	<u>001</u>	Feb 28, 1992
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NORTREL 7/7/7

BARR	<u>0.035MG,0.035MG,0.035MG;0.5MG,0.75MG,1MG</u>	A075478	001	Aug 30, 2002
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TABLET; ORAL-28

ALYACEN 1/35

AB	GLENMARK GENERICS	<u>0.035MG;1MG</u>	<u>A091634</u>	<u>001</u>	Jan 19, 2012
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ALYACEN 7/7/7

AB	GLENMARK GENERICS	<u>0.035MG,0.035MG,0.035MG;0.5MG,0.75MG,1MG</u>	<u>A091636</u>	<u>001</u>	Jan 19, 2012
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ARANELLE

AB	BARR	<u>0.035MG,0.035MG,0.035MG;0.5MG,1MG,0.5MG</u>	<u>A076783</u>	<u>001</u>	Sep 29, 2004
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BALZIVA-28

AB	! BARR	<u>0.035MG;0.4MG</u>	<u>A076238</u>	<u>001</u>	Apr 22, 2004
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BREVICON 28-DAY

AB	ALLERGAN	<u>0.035MG;0.5MG</u>	<u>N017743</u>	<u>001</u>	
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BRIELLYN

AB	GLENMARK GENERICS	<u>0.035MG;0.4MG</u>	<u>A090538</u>	<u>001</u>	Mar 22, 2011
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CYCLAFEM 0.5/35

AB	VINTAGE PHARMS	<u>0.035MG;0.5MG</u>	<u>A203413</u>	<u>001</u>	Dec 16, 2015
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CYCLAFEM 1/35

AB	VINTAGE PHARMS LLC	<u>0.035MG;1MG</u>	<u>A076337</u>	<u>001</u>	Nov 12, 2010
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CYCLAFEM 7/7/7

AB	VINTAGE PHARMS LLC	<u>0.035MG,0.035MG,0.035MG;0.5MG,0.75MG,1MG</u>	<u>A076338</u>	<u>001</u>	Nov 16, 2010
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CYONANZ

AB	AUROBINDO PHARMA LTD	<u>0.035MG;0.5MG</u>	<u>A207055</u>	<u>001</u>	Oct 21, 2016
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DASETTA 1/35

AB	NOVAST LABS LTD	<u>0.035MG;1MG</u>	<u>A090948</u>	<u>001</u>	Dec 22, 2011
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DASETTA 7/7/7

AB	NOVAST LABS LTD	<u>0.035MG,0.035MG,0.035MG;0.5MG,0.75MG,1MG</u>	<u>A090946</u>	<u>001</u>	Dec 22, 2011
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GILDAGIA

AB	VINTAGE PHARMS	<u>0.035MG;0.4MG</u>	<u>A078376</u>	<u>001</u>	Nov 06, 2012
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NORETHINDRONE AND ETHINYL ESTRADIOL

AB	ACCORD HLTHCARE	<u>0.035MG;1MG</u>	<u>A206864</u>	<u>001</u>	Apr 28, 2017
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AB	MAYNE PHARMA	<u>0.035MG;0.5MG</u>	<u>A070686</u>	<u>001</u>	Jan 29, 1987
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AB	WATSON LABS	<u>0.035MG;0.4MG</u>	<u>A078323</u>	<u>001</u>	Feb 04, 2010
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AB	WATSON LABS TEVA	<u>0.035MG;1MG</u>	<u>A070687</u>	<u>001</u>	Jan 29, 1987
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NORINYL 1+35 28-DAY

AB	ALLERGAN	<u>0.035MG;1MG</u>	<u>N017565</u>	<u>002</u>	
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NORTREL 0.5/35-28

AB	BARR	<u>0.035MG;0.5MG</u>	<u>A072695</u>	<u>001</u>	Feb 28, 1992
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PRESCRIPTION DRUG PRODUCT LIST

ETHINYL ESTRADIOL; NORETHINDRONE

TABLET; ORAL-28

NORTREL 1/35-28

AB	! BARR	0.035MG;1MG	A072696 001	Feb 28, 1992
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NORTREL 7/7/7

AB	BARR	0.035MG,0.035MG,0.035MG;0.5MG,0.75MG,1MG	A075478 002	Aug 30, 2002
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NYLIA 1/35

AB	AUROBINDO PHARMA LTD	0.035MG;1MG	A207056 001	Oct 21, 2016
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NYLIA 7/7/7

AB	AUROBINDO PHARMA LTD	0.035MG,0.035MG,0.035MG;0.5MG,0.75MG,1MG	A207054 001	Oct 21, 2016
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ORTHO-NOVUM 7/7/7-28

AB	+ ! JANSSEN PHARMS	0.035MG,0.035MG,0.035MG;0.5MG,0.75MG,1MG	N018985 002	Apr 04, 1984
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PHILITH

AB	NOVAST LABS LTD	0.035MG;0.4MG	A090947 001	Dec 22, 2011
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PIRMELLA 1/35

AB	LUPIN LTD	0.035MG;1MG	A201512 001	Apr 24, 2013
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PIRMELLA 7/7/7

AB	LUPIN LTD	0.035MG,0.035MG,0.035MG;0.5MG,0.75MG,1MG	A201510 001	Apr 24, 2013
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TRI-NORINYL 28-DAY

AB	+ ! MAYNE PHARMA	0.035MG,0.035MG,0.035MG;0.5MG,1MG,0.5MG	N018977 002	Apr 13, 1984
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VYFEMLA

AB	LUPIN LTD	0.035MG;0.4MG	A201886 001	Sep 26, 2013
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WERA

AB	! NOVAST LABS LTD	0.035MG;0.5MG	A091204 001	Mar 27, 2012
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NORETHINDRONE AND ETHINYL ESTRADIOL (10/11)

	WATSON LABS TEVA	0.035MG,0.035MG;0.5MG,1MG	A071044 001	Apr 01, 1988
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TABLET, CHEWABLE; ORAL

KAITLIB FE

AB	LUPIN LTD	0.025MG;0.8MG	A203448 001	Dec 17, 2015
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NEXESTA FE

AB	AUROBINDO PHARMA LTD	0.035MG;0.4MG	A207535 001	Feb 02, 2017
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NORETHINDRONE AND ETHINYL ESTRADIOL AND FERROUS FUMARATE

AB	ACCORD HLTHCARE	0.035MG;0.4MG	A207066 001	Mar 29, 2017
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AB	AMNEAL PHARMS	0.035MG;0.4MG	A078892 001	Sep 26, 2011
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AB	+ ! APIL	0.025MG;0.8MG	N022573 001	Dec 22, 2010
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AB	BARR	0.035MG;0.4MG	A078965 001	Aug 05, 2010
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AB	LUPIN LTD	0.035MG;0.4MG	A091332 001	Mar 23, 2016
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AB	MYLAN LABS LTD	0.025MG;0.8MG	A203371 001	Apr 23, 2014
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AB	!	0.035MG;0.4MG	A202086 001	Apr 01, 2015
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ETHINYL ESTRADIOL; NORETHINDRONE ACETATE

CAPSULE; ORAL

TAYTULLA

	+ ! APIL	0.02MG;1MG	N204426 001	Apr 19, 2013
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TABLET; ORAL

AUROVELA 24 FE

AB	AUROBINDO PHARMA LTD	0.02MG;1MG	A207504 001	Jun 15, 2017
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BLISOVI 24 FE

AB	LUPIN LTD	0.02MG;1MG	A091398 001	Oct 28, 2015
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FEMHRT

AB	APIL	0.0025MG;0.5MG	N021065 001	Jan 14, 2005
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FYAVOLV

AB	LUPIN LTD	0.005MG;1MG	A204213 002	Dec 10, 2015
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AB		0.0025MG;0.5MG	A204213 001	Dec 10, 2015
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GILDESS 24 FE

AB	VINTAGE PHARMS	0.02MG;1MG	A090293 001	Dec 01, 2014
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LARIN 24 FE

AB	NOVAST LABS	0.02MG;1MG	A202994 001	Feb 18, 2015
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LERIBANE

AB	NOVAST LABS	0.0025MG;0.5MG	A203435 002	Jun 03, 2016
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AB		0.005MG;1MG	A203435 001	Jun 03, 2016
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LOESTRIN 24 FE

AB	+ TEVA BRANDED PHARM	0.02MG;1MG	N021871 001	Feb 17, 2006
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NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL

AB	! BARR LABS INC	0.005MG;1MG	A076221 001	Nov 06, 2009
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AB	GLENMARK GENERICS	0.0025MG;0.5MG	A203038 001	Apr 02, 2015
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AB		0.005MG;1MG	A203038 002	Apr 02, 2015
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PRESCRIPTION DRUG PRODUCT LIST

ETHINYL ESTRADIOL; NORETHINDRONE ACETATE

TABLET; ORAL

NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL

AB	MYLAN LABS LTD	<u>0.0025MG;0.5MG</u>	<u>A207260 001</u>	Feb 02, 2017
AB		<u>0.005MG;1MG</u>	<u>A207259 001</u>	Dec 27, 2016

NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL AND FERROUS FUMARATE

AB	! AMNEAL PHARMS	<u>0.02MG;1MG</u>	<u>A078267 001</u>	Sep 01, 2009
AB	BARR LABS INC	<u>0.02MG;1MG</u>	<u>A090938 001</u>	Dec 01, 2014
AB	GLENMARK PHARMS LTD	<u>0.02MG;1MG</u>	<u>A204847 001</u>	Nov 17, 2017
AB	MYLAN LABS LTD	<u>0.02MG;1MG</u>	<u>A202742 001</u>	Oct 30, 2014
	LO LOESTRIN FE			
	+! APIL	0.01MG,0.01MG;1MG,N/A	N022501 001	Oct 21, 2010

TABLET; ORAL-21

AUROVELA 1.5/30

AB	AUROBINDO PHARMA LTD	<u>0.03MG;1.5MG</u>	<u>A207581 001</u>	Jun 26, 2017
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AUROVELA 1/20

AB	AUROBINDO PHARMA LTD	<u>0.02MG;1MG</u>	<u>A207506 001</u>	Jun 16, 2017
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GILDESS 1.5/30

AB	VINTAGE PHARMS LLC	<u>0.03MG;1.5MG</u>	<u>A077075 002</u>	Jul 24, 2012
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GILDESS 1/20

AB	VINTAGE PHARMS LLC	<u>0.02MG;1MG</u>	<u>A077077 002</u>	Jul 24, 2012
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HAILEY 1.5/30

AB	GLENMARK PHARMS	<u>0.03MG;1.5MG</u>	<u>A209297 001</u>	Jun 05, 2018
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JUNEL 1.5/30

AB	BARR	<u>0.03MG;1.5MG</u>	<u>A076381 001</u>	May 30, 2003
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JUNEL 1/20

AB	BARR	<u>0.02MG;1MG</u>	<u>A076380 001</u>	May 30, 2003
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LARIN 1.5/30

AB	NOVAST LABS	<u>0.03MG;1.5MG</u>	<u>A202996 001</u>	Mar 20, 2014
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LARIN 1/20

AB	NOVAST LABS	<u>0.02MG;1MG</u>	<u>A202995 001</u>	Dec 04, 2013
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LOESTRIN 21 1.5/30

AB	+ TEVA BRANDED PHARM	<u>0.03MG;1.5MG</u>	<u>N017875 001</u>	
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LOESTRIN 21 1/20

AB	+ TEVA BRANDED PHARM	<u>0.02MG;1MG</u>	<u>N017876 001</u>	
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LOESTRIN FE 1.5/30

AB	+! TEVA BRANDED PHARM	<u>0.03MG;1.5MG</u>	<u>N017355 001</u>	
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MICROGESTIN 1.5/30

AB	MAYNE PHARMA	<u>0.03MG;1.5MG</u>	<u>A075548 002</u>	Jul 30, 2003
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MICROGESTIN 1/20

AB	MAYNE PHARMA	<u>0.02MG;1MG</u>	<u>A075647 002</u>	Jul 30, 2003
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NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL

AB	GLENMARK PHARMS LTD	<u>0.02MG;1MG</u>	<u>A206969 001</u>	Jan 20, 2016
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AB	MYLAN LABS LTD	<u>0.02MG;1MG</u>	<u>A202771 001</u>	Nov 06, 2013
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AB		<u>0.03MG;1.5MG</u>	<u>A202770 001</u>	Feb 19, 2015
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TRI-LEGEST 21

BARR 0.02MG,0.03MG,0.035MG;1MG,1MG,1MG

A076405 001 Oct 26, 2007

TABLET; ORAL-28

AUROVELA FE 1.5/30

AB	AUROBINDO PHARMA LTD	<u>0.03MG;1.5MG</u>	<u>A207580 001</u>	Jun 15, 2017
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AUROVELA FE 1/20

AB	AUROBINDO PHARMA LTD	<u>0.02MG;1MG</u>	<u>A207505 001</u>	Jun 16, 2017
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BLISOVI FE 1.5/30

AB	LUPIN LTD	<u>0.03MG;1.5MG</u>	<u>A201585 001</u>	Nov 18, 2015
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BLISOVI FE 1/20

AB	LUPIN LTD	<u>0.02MG;1MG</u>	<u>A201584 001</u>	Nov 18, 2015
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ESTROSTEP FE

AB	+! APIL	<u>0.02MG,0.03MG,0.035MG;1MG,1MG,1MG</u>	<u>N020130 002</u>	Oct 09, 1996
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GILDESS FE 1.5/30

AB	VINTAGE PHARMS LLC	<u>0.03MG;1.5MG</u>	<u>A077075 001</u>	Apr 28, 2005
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GILDESS FE 1/20

AB	VINTAGE PHARMS LLC	<u>0.02MG;1MG</u>	<u>A077077 001</u>	May 20, 2005
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HAILEY FE 1.5/30

AB	GLENMARK PHARMS	<u>0.03MG;1.5MG</u>	<u>A209031 001</u>	Jun 05, 2018
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HAILEY FE 1/20

AB	GLENMARK PHARMS LTD	<u>0.02MG;1MG</u>	<u>A206597 001</u>	Nov 21, 2017
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JUNEL FE 1.5/30

AB	BARR	<u>0.03MG;1.5MG</u>	<u>A076064 001</u>	Sep 18, 2003
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PRESCRIPTION DRUG PRODUCT LIST

ETHINYL ESTRADIOL; NORETHINDRONE ACETATE

TABLET; ORAL-28

JUNEL FE 1/20

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>LOESTRIN FE 1/20</u>			
AB	+ TEVA BRANDED PHARM	0.02MG;1MG	N017354 001	
	<u>MICROGESTIN FE 1.5/30</u>			
AB	MAYNE PHARMA	0.03MG;1.5MG	A075548 001	Feb 05, 2001
	<u>MICROGESTIN FE 1/20</u>			
AB	MAYNE PHARMA	0.02MG;1MG	A075647 001	Feb 05, 2001
	<u>NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL</u>			
AB	MAYNE PHARMA	0.02MG,0.03MG,0.035MG;1MG,1MG,1MG	A076629 001	Mar 18, 2010
AB	MYLAN LABS LTD	0.02MG;1MG	A202772 001	Nov 14, 2013
AB		0.03MG;1.5MG	A202741 001	Feb 20, 2015
	<u>TRI-LEGEST FE</u>			
AB	BARR	0.02MG,0.03MG,0.035MG;1MG,1MG,1MG	A076105 001	Oct 26, 2007
	TABLET, CHEWABLE; ORAL			
	<u>MIBELAS 24 FE</u>			
AB	LUPIN ATLANTIS	0.02MG;1MG	A206287 001	May 24, 2016
	<u>MINASTRIN 24 FE</u>			
AB	+! APIL	0.02MG;1MG	N203667 001	May 08, 2013
	<u>NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL AND FERROUS FUMARATE</u>			
AB	AMNEAL PHARMS	0.02MG;1MG	A207514 001	Sep 11, 2017
AB	GLENMARK PHARMS LTD	0.02MG;1MG	A210369 001	Dec 26, 2017
AB	XIROMED	0.02MG;1MG	A209609 001	Jul 16, 2018

ETHINYL ESTRADIOL; NORGESTIMATE

TABLET; ORAL-28

ESTARYLLA

AB	XIROMED	0.035MG;0.25MG	A090794 001	Jan 30, 2013
	<u>MILI</u>			
AB	AUROBINDO PHARMA LTD	0.035MG;0.25MG	A205449 001	Jul 07, 2016
	<u>MONO-LINYAH</u>			
AB	NOVAST LABS LTD	0.035MG;0.25MG	A090523 001	May 23, 2012
	<u>NORGESTIMATE AND ETHINYL ESTRADIOL</u>			
AB	AMNEAL PHARMS	0.035MG,0.035MG,0.035MG;0.18MG,0.215MG,0.25MG	A203870 001	Nov 12, 2015
AB		0.035MG;0.25MG	A203865 001	Oct 27, 2015
AB		0.025MG,0.025MG,0.025MG;0.18MG,0.215MG,0.25MG	A203873 001	May 12, 2016
AB	GLENMARK GENERICS	0.035MG,0.035MG,0.035MG;0.18MG,0.215MG,0.25MG	A200494 001	Jun 17, 2011
AB		0.035MG;0.25MG	A200538 001	Apr 05, 2012
AB	GLENMARK PHARMS	0.025MG,0.025MG,0.025MG;0.18MG,0.215MG,0.25MG	A204057 001	Feb 23, 2016
AB	LUPIN LTD	0.035MG,0.035MG,0.035MG;0.18MG,0.215MG,0.25MG	A205588 001	Apr 26, 2016
AB		0.035MG;0.25MG	A205630 001	Oct 27, 2016
AB	LUPIN PHARMS	0.025MG,0.025MG,0.025MG;0.18MG,0.215MG,0.25MG	A200541 001	Jun 25, 2012
AB	MYLAN LABS LTD	0.035MG,0.035MG,0.035MG;0.18MG,0.215MG,0.25MG	A201897 001	Jan 27, 2016
AB		0.035MG;0.25MG	A201896 001	Jan 27, 2016
AB	NAARI PTE LTD	0.035MG;0.035MG;0.035MG;0.18MG;0.215MG;0.25MG	A200383 001	Apr 07, 2015
AB		0.035MG;0.25MG	A200384 001	Apr 07, 2015
	<u>ORTHO TRI-CYCLEN</u>			
AB	+! JANSSEN PHARMS	0.035MG,0.035MG,0.035MG;0.18MG,0.215MG,0.25MG	N019697 001	Jul 03, 1992
	<u>ORTHO TRI-CYCLEN LO</u>			
AB	+! JANSSEN PHARMS	0.025MG,0.025MG,0.025MG;0.18MG,0.215MG,0.25MG	N021241 001	Aug 22, 2002
	<u>PREVIFEM</u>			
AB	VINTAGE PHARMS LLC	0.035MG;0.25MG	A076334 001	Jan 09, 2004
	<u>SPRINTEC</u>			
AB	! BARR	0.035MG;0.25MG	A075804 001	Sep 25, 2002
	<u>TRI LO SPRINTEC</u>			
AB	BARR LABS INC	0.025MG,0.025MG,0.025MG;0.18MG,0.215MG,0.25MG	A076784 001	Jun 29, 2009

PRESCRIPTION DRUG PRODUCT LIST

ETHINYL ESTRADIOL; NORGESTIMATE

TABLET; ORAL-28

TRI-ESTARYLLA

AB	XIROMED	<u>0.035MG, 0.035MG, 0.035MG; 0.18MG, 0.215MG, 0.25MG</u>	<u>A090793 001</u>	Jan 30, 2013
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TRI-LINYAH

AB	NOVAST LABS LTD	<u>0.035MG, 0.035MG, 0.035MG; 0.18MG, 0.215MG, 0.25MG</u>	<u>A090524 001</u>	May 30, 2012
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TRI-LO-ESTARYLLA

AB	XIROMED	<u>0.025MG, 0.025MG, 0.025MG; 0.18MG, 0.215MG, 0.25MG</u>	<u>A091232 001</u>	Jun 29, 2015
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TRI-LO-MILI

AB	AUROBINDO PHARMA LTD	<u>0.025MG, 0.025MG, 0.025MG; 0.18MG, 0.215MG, 0.25MG</u>	<u>A205762 001</u>	Nov 04, 2016
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TRI-MILI

AB	AUROBINDO PHARMA LTD	<u>0.035MG, 0.035MG, 0.035MG; 0.18MG, 0.215MG, 0.25MG</u>	<u>A205441 001</u>	Jul 06, 2016
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TRI-PREVI-FEM

AB	VINTAGE PHARMS LLC	<u>0.035MG, 0.035MG, 0.035MG; 0.18MG, 0.215MG, 0.25MG</u>	<u>A076335 001</u>	Mar 26, 2004
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TRI-SPRINTEC

AB	BARR	<u>0.035MG, 0.035MG, 0.035MG; 0.18MG, 0.215MG, 0.25MG</u>	<u>A075808 001</u>	Dec 29, 2003
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ETHINYL ESTRADIOL; NORGESTREL

TABLET; ORAL-21

CRYSSELLE

AB	DURAMED PHARMS BARR	<u>0.03MG; 0.3MG</u>	<u>A075840 001</u>	Nov 30, 2001
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TABLET; ORAL-28

CRYSSELLE

AB	DURAMED PHARMS BARR	<u>0.03MG; 0.3MG</u>	<u>A075840 002</u>	Nov 30, 2001
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ELINEST

AB	NOVAST LABS LTD	<u>0.03MG; 0.3MG</u>	<u>A091105 001</u>	Mar 28, 2012
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LOW-OGESTREL-28

AB	MAYNE PHARMA	<u>0.03MG; 0.3MG</u>	<u>A075288 002</u>	Jul 28, 1999
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OGESTREL 0.5/50-28

!	WATSON LABS	0.05MG; 0.5MG	A075406 002	Dec 15, 1999
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ETHINYL ESTRADIOL; SEGESTERONE ACETATE

RING; VAGINAL

ANNOVERA

+	THERAPEUTICSMD INC	0.013MG/24HR; 0.15MG/24HR	N209627 001	Aug 10, 2018
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ETHIODIZED OIL

OIL; INTRALYMPHATIC, INTRAUTERINE

LIPIODOL

+	GUERBET	EQ 4.8GM IODINE/10ML (EQ 480MG IODINE/ML)	N009190 001	
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ETHIONAMIDE

TABLET; ORAL

TRECATOR

+	WYETH PHARMS	250MG	N013026 002	
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ETHOSUXIMIDE

CAPSULE; ORAL

ETHOSUXIMIDE

AB	AKORN	<u>250MG</u>	<u>A040686 001</u>	May 28, 2008
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AB	BIONPHARMA INC	<u>250MG</u>	<u>A040430 001</u>	Oct 28, 2002
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AB	HERITAGE PHARMS INC	<u>250MG</u>	<u>A200892 001</u>	Sep 25, 2012
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AB	PURACAP PHARM LLC	<u>250MG</u>	<u>A210654 001</u>	Mar 16, 2020
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AB	STRIDES PHARMA	<u>250MG</u>	<u>A211928 001</u>	Feb 19, 2019
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ZARONTIN

AB	+	PARKE DAVIS	<u>250MG</u>	<u>N012380 001</u>
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SYRUP; ORAL

ETHOSUXIMIDE

AA	MIKART	<u>250MG/5ML</u>	<u>A040506 001</u>	Dec 22, 2003
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AA	PHARM ASSOC	<u>250MG/5ML</u>	<u>A040253 001</u>	Nov 22, 2000
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ZARONTIN

AA	!	PARKE-DAVIS	<u>250MG/5ML</u>	<u>A080258 001</u>
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PRESCRIPTION DRUG PRODUCT LIST

ETHOTOIN

TABLET; ORAL

PEGANONE

+! RECORDATI RARE 250MG N010841 001

ETODOLAC

CAPSULE; ORAL

ETODOLAC

<u>AB</u>	ANI PHARMS INC	<u>200MG</u>	<u>A075126 001</u>	Sep 16, 1999
<u>AB</u>		<u>300MG</u>	<u>A075126 002</u>	Sep 16, 1999
<u>AB</u>	APOTEX	<u>200MG</u>	<u>A075419 001</u>	Jul 28, 2000
<u>AB</u>		<u>300MG</u>	<u>A075419 002</u>	Jul 28, 2000
<u>AB</u>	TARO	<u>200MG</u>	<u>A075078 001</u>	Apr 30, 1998
<u>AB</u>	!	<u>300MG</u>	<u>A075078 002</u>	Apr 30, 1998

TABLET; ORAL

ETODOLAC

<u>AB</u>	AMNEAL PHARMS CO	<u>400MG</u>	<u>A208834 001</u>	Jun 07, 2018
<u>AB</u>		<u>500MG</u>	<u>A208834 002</u>	Jun 07, 2018
<u>AB</u>	APOTEX INC	<u>400MG</u>	<u>A076004 001</u>	Dec 03, 2002
<u>AB</u>		<u>500MG</u>	<u>A076004 002</u>	Dec 03, 2002
<u>AB</u>	EDENBRIDGE PHARMS	<u>400MG</u>	<u>A209888 001</u>	Nov 30, 2018
<u>AB</u>		<u>500MG</u>	<u>A209888 002</u>	Nov 30, 2018
<u>AB</u>	SANDOZ	<u>400MG</u>	<u>A074903 001</u>	Apr 11, 1997
<u>AB</u>		<u>500MG</u>	<u>A074903 002</u>	Apr 19, 1999
<u>AB</u>	TARO PHARM INDS	<u>400MG</u>	<u>A075074 001</u>	Mar 11, 1998
<u>AB</u>	!	<u>500MG</u>	<u>A075074 002</u>	Apr 25, 2000

TABLET, EXTENDED RELEASE; ORAL

ETODOLAC

<u>AB</u>	TARO	<u>400MG</u>	<u>A076174 001</u>	Mar 13, 2003
<u>AB</u>		<u>500MG</u>	<u>A076174 002</u>	Mar 13, 2003
<u>AB</u>		<u>600MG</u>	<u>A076174 003</u>	Mar 13, 2003
<u>AB</u>	TEVA	<u>400MG</u>	<u>A075665 003</u>	Feb 05, 2001
<u>AB</u>		<u>500MG</u>	<u>A075665 002</u>	Jul 31, 2000
<u>AB</u>	!	<u>600MG</u>	<u>A075665 001</u>	Jul 31, 2000
<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
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ETOMIDATE

<u>AP</u>	AUROBINDO PHARMA LTD	<u>2MG/ML</u>	<u>A206126 001</u>	Feb 24, 2017
<u>AP</u>	EMCURE PHARMS LTD	<u>2MG/ML</u>	<u>A204618 001</u>	Aug 13, 2014
<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>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>	WEST-WARD PHARMS INT	<u>2MG/ML</u>	<u>A074593 001</u>	Nov 04, 1996
<u>AP</u>	ZYDUS PHARMS	<u>2MG/ML</u>	<u>A202360 001</u>	Jul 18, 2014

ETONOGESTREL

IMPLANT; IMPLANTATION

NEXPLANON

+! ORGANON USA INC 68MG/IMPLANT N021529 002 May 31, 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>	MYLAN LABS LTD	<u>20MG/ML</u>	<u>A203507 001</u>	Nov 20, 2017
<u>AP</u>		<u>20MG/ML</u>	<u>A204927 001</u>	Oct 31, 2017
<u>AP</u>	TEVA PHARMS USA	<u>20MG/ML</u>	<u>A074529 001</u>	Jul 24, 1996
<u>AP</u>	WEST-WARD PHARMS INT	<u>20MG/ML</u>	<u>A074290 001</u>	Jul 17, 1995

PRESCRIPTION DRUG PRODUCT LIST

ETOPOSID PHOSPHATE

INJECTABLE; INJECTION

ETOPOPHOS PRESERVATIVE FREE

+! CHEPLAPHARM EQ 100MG BASE/VIAL N020457 001 May 17, 1996

ETRAVIRINE

TABLET; ORAL

INTELENCE

+ JANSSEN R AND D 25MG N022187 003 Mar 26, 2012

+ 100MG N022187 001 Jan 18, 2008

+! 200MG N022187 002 Dec 22, 2010

EVEROLIMUS

TABLET; ORAL

AFINITORAB + NOVARTIS 2.5MG N022334 003 Jul 09, 2010AB +! 5MG N022334 001 Mar 30, 2009AB + 7.5MG N022334 004 Mar 30, 2012AB + 10MG N022334 002 Mar 30, 2009EVEROLIMUSAB HIKMA 0.25MG A206133 001 Apr 12, 2018AB 0.5MG A206133 002 Apr 12, 2018AB 0.75MG A206133 003 Apr 12, 2018AB PAR PHARM 2.5MG A207934 001 Dec 09, 2019AB 5MG A207934 002 Dec 09, 2019AB 7.5MG A207934 003 Dec 09, 2019AB TEVA PHARMS USA 2.5MG A210050 001 Dec 09, 2019AB 5MG A210050 002 Dec 09, 2019AB 7.5MG A210050 003 Dec 09, 2019AB 10MG A210050 004 Dec 09, 2019ZORTRESSAB + NOVARTIS 0.25MG N021560 001 Apr 20, 2010AB + 0.5MG N021560 002 Apr 20, 2010AB +! 0.75MG N021560 003 Apr 20, 2010

+ 1MG N021560 004 Aug 10, 2018

TABLET, FOR SUSPENSION; ORAL

AFINITOR DISPERZ

+ NOVARTIS PHARM 2MG N203985 001 Aug 29, 2012

+ 3MG N203985 002 Aug 29, 2012

+! 5MG N203985 003 Aug 29, 2012

EXEMESTANE

TABLET; ORAL

AROMASINAB +! PHARMACIA AND UPJOHN 25MG N020753 001 Oct 21, 1999EXEMESTANEAB ALVOGEN 25MG A200898 001 Jul 28, 2014AB BRECKENRIDGE 25MG A211031 001 Feb 21, 2019AB CIPLA 25MG A210323 001 Apr 27, 2018AB HIKMA 25MG A077431 001 Apr 01, 2011AB MYLAN 25MG A203315 001 Mar 10, 2017AB UPSHER SMITH LABS 25MG A209208 001 Jul 26, 2017AB ZYDUS PHARMS 25MG A202602 001 Oct 03, 2018EXENATIDE

SUSPENSION, EXTENDED RELEASE; SUBCUTANEOUS

BYDUREON BCISE

+! ASTRAZENECA AB 2MG/0.85ML (2MG/0.85ML) N209210 001 Oct 20, 2017

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

INJECTABLE; SUBCUTANEOUS

BYETTA

+! ASTRAZENECA AB 300MCG/1.2ML (250MCG/ML) N021773 001 Apr 28, 2005

+! 600MCG/2.4ML (250MCG/ML) N021773 002 Apr 28, 2005

PRESCRIPTION DRUG PRODUCT LIST

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>	APOTEX	<u>10MG</u>	<u>A208332</u>	<u>001</u>	Jun 12, 2017
<u>AB</u>	AUROBINDO PHARMA LTD	<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>	OHM LABS INC	<u>10MG</u>	<u>A207311</u>	<u>001</u>	Jun 12, 2017
<u>AB</u>	SANDOZ INC	<u>10MG</u>	<u>A203931</u>	<u>001</u>	Jun 12, 2017
<u>AB</u>	TEVA PHARMS USA	<u>10MG</u>	<u>A078724</u>	<u>001</u>	Jun 12, 2017
<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>	<u>+</u> !	MSD INTL GMBH	<u>10MG</u>	<u>N021445</u>	<u>001</u>	Oct 25, 2002
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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>	ANI PHARMS INC	<u>10MG; 10MG</u>	<u>A201890</u>	<u>001</u>	Apr 26, 2017
<u>AB</u>		<u>10MG; 20MG</u>	<u>A201890</u>	<u>002</u>	Apr 26, 2017
<u>AB</u>		<u>10MG; 40MG</u>	<u>A201890</u>	<u>003</u>	Apr 26, 2017
<u>AB</u>		<u>10MG; 80MG</u>	<u>A201890</u>	<u>004</u>	Apr 26, 2017
<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>	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>	<u>+</u>	MSD INTL	<u>10MG; 10MG</u>	<u>N021687</u>	<u>001</u>	Jul 23, 2004
<u>AB</u>	<u>+</u>		<u>10MG; 20MG</u>	<u>N021687</u>	<u>002</u>	Jul 23, 2004
<u>AB</u>	<u>+</u>		<u>10MG; 40MG</u>	<u>N021687</u>	<u>003</u>	Jul 23, 2004
<u>AB</u>	<u>+</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>	CIPLA	<u>125MG</u>	<u>A078278</u>	<u>001</u>	Mar 21, 2011
<u>AB</u>		<u>250MG</u>	<u>A078278</u>	<u>002</u>	Mar 21, 2011
<u>AB</u>		<u>500MG</u>	<u>A078278</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
<u>AB</u>		<u>500MG</u>	<u>A202438</u>	<u>003</u>	Sep 10, 2014
<u>AB</u>	HIKMA	<u>125MG</u>	<u>A090128</u>	<u>001</u>	Mar 21, 2011
<u>AB</u>		<u>250MG</u>	<u>A090128</u>	<u>002</u>	Mar 21, 2011
<u>AB</u>		<u>500MG</u>	<u>A090128</u>	<u>003</u>	Mar 21, 2011
<u>AB</u>	MACLEODS PHARMS LTD	<u>125MG</u>	<u>A201022</u>	<u>001</u>	Jan 12, 2012
<u>AB</u>		<u>250MG</u>	<u>A201022</u>	<u>002</u>	Jan 12, 2012
<u>AB</u>		<u>500MG</u>	<u>A201022</u>	<u>003</u>	Jan 12, 2012
<u>AB</u>	MYLAN	<u>125MG</u>	<u>A201333</u>	<u>001</u>	Mar 24, 2011

PRESCRIPTION DRUG PRODUCT LIST

FAMCICLOVIR

TABLET; ORAL

FAMCICLOVIR

<u>AB</u>		<u>250MG</u>	<u>A201333 002</u>	Mar 24, 2011
<u>AB</u>		<u>500MG</u>	<u>A201333 003</u>	Mar 24, 2011
<u>AB</u>	TEVA PHARMS	<u>125MG</u>	<u>A077487 001</u>	Aug 24, 2007
<u>AB</u>		<u>250MG</u>	<u>A077487 002</u>	Aug 24, 2007
<u>AB</u>	!	<u>500MG</u>	<u>A077487 003</u>	Aug 24, 2007

FAMOTIDINE

FOR SUSPENSION; ORAL

FAMOTIDINE

<u>AB</u>	HI-TECH PHARMA CO	<u>40MG/5ML</u>	<u>A201995 001</u>	May 30, 2014
<u>AB</u>	!	<u>40MG/5ML</u>	<u>A090440 001</u>	Jun 29, 2010
<u>AB</u>	NAVINTA LLC	<u>40MG/5ML</u>	<u>A091020 001</u>	May 27, 2010
<u>AB</u>	NOVEL LABS INC	<u>40MG/5ML</u>	<u>A201695 001</u>	Dec 17, 2012

INJECTABLE; INJECTION

FAMOTIDINE

<u>AP</u>	ATHENEX INC	<u>10MG/ML</u>	<u>A075651 001</u>	Apr 16, 2001
<u>AP</u>		<u>10MG/ML</u>	<u>A075684 001</u>	Apr 16, 2001
<u>AP</u>	FRESENIUS KABI USA	<u>10MG/ML</u>	<u>A075709 001</u>	Apr 16, 2001
<u>AP</u>	MYLAN LABS LTD	<u>10MG/ML</u>	<u>A078641 001</u>	Jun 25, 2008
<u>AP</u>	!	<u>10MG/ML</u>	<u>A075488 001</u>	Apr 16, 2001

FAMOTIDINE PRESERVATIVE FREE

<u>AP</u>	ATHENEX INC	<u>10MG/ML</u>	<u>A075622 001</u>	Apr 16, 2001
<u>AP</u>		<u>10MG/ML</u>	<u>A075825 001</u>	Apr 17, 2001
<u>AP</u>	FRESENIUS KABI USA	<u>10MG/ML</u>	<u>A075813 001</u>	Apr 16, 2001
<u>AP</u>	MYLAN LABS LTD	<u>10MG/ML</u>	<u>A078642 001</u>	Jun 25, 2008
<u>AP</u>	!	<u>10MG/ML</u>	<u>A075486 001</u>	Apr 16, 2001

INT

FAMOTIDINE PRESERVATIVE FREE IN PLASTIC CONTAINER

! BAXTER HLTHCARE 0.4MG/ML

A075591 001 May 10, 2001

TABLET; ORAL

FAMOTIDINE

<u>AB</u>	ALEMBIC PHARMS LTD	<u>20MG</u>	<u>A078916 001</u>	May 22, 2009
<u>AB</u>		<u>40MG</u>	<u>A078916 002</u>	May 22, 2009
<u>AB</u>	APOTEX	<u>20MG</u>	<u>A075611 001</u>	Jul 23, 2001
<u>AB</u>		<u>40MG</u>	<u>A075611 002</u>	Jul 23, 2001
<u>AB</u>	AUROBINDO PHARMA LTD	<u>20MG</u>	<u>A206530 001</u>	Dec 22, 2015
<u>AB</u>	!	<u>40MG</u>	<u>A206530 002</u>	Dec 22, 2015
<u>AB</u>	CARLSBAD	<u>20MG</u>	<u>A075805 001</u>	Apr 16, 2001
<u>AB</u>		<u>40MG</u>	<u>A075805 002</u>	Apr 16, 2001
<u>AB</u>	CELLTRION	<u>20MG</u>	<u>A075786 001</u>	Apr 16, 2001
<u>AB</u>		<u>40MG</u>	<u>A075786 002</u>	Apr 16, 2001
<u>AB</u>	DR REDDYS LABS LTD	<u>20MG</u>	<u>A075718 001</u>	Apr 16, 2001
<u>AB</u>		<u>40MG</u>	<u>A075718 002</u>	Apr 16, 2001
<u>AB</u>	IVAX SUB TEVA PHARMS	<u>20MG</u>	<u>A075511 001</u>	Apr 16, 2001
<u>AB</u>		<u>40MG</u>	<u>A075511 002</u>	Apr 16, 2001
<u>AB</u>	PERRIGO R AND D	<u>20MG</u>	<u>A077352 002</u>	Jul 27, 2005
<u>AB</u>		<u>40MG</u>	<u>A077352 001</u>	Jul 27, 2005
<u>AB</u>	TEVA	<u>20MG</u>	<u>A075311 001</u>	Apr 16, 2001
<u>AB</u>		<u>40MG</u>	<u>A075311 002</u>	Apr 16, 2001

FAMOTIDINE; IBUPROFEN

TABLET; ORAL

DUEXIS

+! HORIZON 26.6MG;800MG

N022519 001 Apr 23, 2011

FEBUXOSTAT

TABLET; ORAL

FEBUXOSTAT

<u>AB</u>	ALEMBIC PHARMS LTD	<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>	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 REMEDIES	<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>	MSN	<u>40MG</u>	<u>A210461 001</u>	Dec 30, 2019
<u>AB</u>		<u>80MG</u>	<u>A210461 002</u>	Dec 30, 2019
<u>AB</u>	MYLAN	<u>40MG</u>	<u>A205385 001</u>	Jul 01, 2019
<u>AB</u>		<u>80MG</u>	<u>A205385 002</u>	Jul 01, 2019
<u>AB</u>	SUN PHARM	<u>40MG</u>	<u>A205467 001</u>	Jul 01, 2019

PRESCRIPTION DRUG PRODUCT LIST

FEBUXOSTAT

TABLET; ORAL

FEBUXOSTAT

<u>AB</u>		<u>80MG</u>	<u>A205467 002</u>	Jul 01, 2019
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ULORIC

<u>AB</u>	+	TAKEDA PHARMS USA	<u>40MG</u>	<u>N021856 001</u>	Feb 13, 2009
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<u>AB</u>	+	!	<u>80MG</u>	<u>N021856 002</u>	Feb 13, 2009
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FEDRATINIB HYDROCHLORIDE

CAPSULE; ORAL

INREBIC

+	!	IMPACT	EQ 100MG BASE	N212327 001	Aug 16, 2019
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FELBAMATE

SUSPENSION; ORAL

FELBAMATE

<u>AB</u>		AMNEAL PHARMS	<u>600MG/5ML</u>	<u>A202385 001</u>	Dec 16, 2011
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<u>AB</u>		TARO	<u>600MG/5ML</u>	<u>A206314 001</u>	Jun 16, 2017
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<u>AB</u>		VISTAPHARM	<u>600MG/5ML</u>	<u>A211333 001</u>	May 31, 2019
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FELBATOL

<u>AB</u>	+	!	MYLAN SPECIALITY LP	<u>600MG/5ML</u>	<u>N020189 003</u>	Jul 29, 1993
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TABLET; ORAL

FELBAMATE

<u>AB</u>		ALVOGEN	<u>400MG</u>	<u>A204595 001</u>	Jan 11, 2016
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<u>AB</u>			<u>600MG</u>	<u>A204595 002</u>	Jan 11, 2016
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<u>AB</u>		AMNEAL PHARMS	<u>400MG</u>	<u>A201680 001</u>	Sep 13, 2011
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<u>AB</u>			<u>600MG</u>	<u>A201680 002</u>	Sep 13, 2011
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<u>AB</u>		ANI PHARMS INC	<u>400MG</u>	<u>A202284 001</u>	Nov 04, 2015
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<u>AB</u>			<u>600MG</u>	<u>A202284 002</u>	Nov 04, 2015
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<u>AB</u>		CADILA	<u>400MG</u>	<u>A208970 001</u>	May 30, 2017
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<u>AB</u>			<u>600MG</u>	<u>A208970 002</u>	May 30, 2017
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<u>AB</u>		TARO	<u>400MG</u>	<u>A207093 001</u>	Apr 20, 2017
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<u>AB</u>			<u>600MG</u>	<u>A207093 002</u>	Apr 20, 2017
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FELBATOL

<u>AB</u>	+		MYLAN SPECIALITY LP	<u>400MG</u>	<u>N020189 001</u>	Jul 29, 1993
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<u>AB</u>	+	!		<u>600MG</u>	<u>N020189 002</u>	Jul 29, 1993
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FELODIPINE

TABLET, EXTENDED RELEASE; ORAL

FELODIPINE

<u>AB</u>		AUROBINDO PHARMA LTD	<u>2.5MG</u>	<u>A203417 001</u>	Jan 17, 2013
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<u>AB</u>			<u>5MG</u>	<u>A203417 002</u>	Jan 17, 2013
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<u>AB</u>			<u>10MG</u>	<u>A203417 003</u>	Jan 17, 2013
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<u>AB</u>		GLENMARK GENERICS	<u>2.5MG</u>	<u>A090365 001</u>	Dec 17, 2010
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<u>AB</u>			<u>5MG</u>	<u>A090365 002</u>	Dec 17, 2010
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<u>AB</u>			<u>10MG</u>	<u>A090365 003</u>	Dec 17, 2010
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<u>AB</u>		HERITAGE PHARMS INC	<u>2.5MG</u>	<u>A201964 001</u>	Nov 08, 2013
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<u>AB</u>			<u>5MG</u>	<u>A201964 002</u>	Nov 08, 2013
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<u>AB</u>			<u>10MG</u>	<u>A201964 003</u>	Nov 08, 2013
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<u>AB</u>		JUBILANT GENERICS	<u>2.5MG</u>	<u>A203983 001</u>	Aug 19, 2016
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<u>AB</u>			<u>5MG</u>	<u>A203983 002</u>	Aug 19, 2016
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<u>AB</u>			<u>10MG</u>	<u>A203983 003</u>	Aug 19, 2016
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<u>AB</u>		ORCHID HLTHCARE	<u>2.5MG</u>	<u>A203032 001</u>	May 21, 2015
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<u>AB</u>			<u>5MG</u>	<u>A203032 002</u>	May 21, 2015
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<u>AB</u>			<u>10MG</u>	<u>A203032 003</u>	May 21, 2015
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<u>AB</u>		SUN PHARM INDS LTD	<u>2.5MG</u>	<u>A091200 001</u>	Dec 13, 2013
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<u>AB</u>			<u>5MG</u>	<u>A091200 002</u>	Dec 13, 2013
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<u>AB</u>			<u>10MG</u>	<u>A091200 003</u>	Dec 13, 2013
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<u>AB</u>		SUN PHARM INDUSTRIES	<u>2.5MG</u>	<u>A075896 001</u>	Nov 02, 2004
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<u>AB</u>			<u>5MG</u>	<u>A075896 002</u>	Nov 02, 2004
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<u>AB</u>			<u>10MG</u>	<u>A075896 003</u>	Nov 02, 2004
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<u>AB</u>		TORRENT PHARMS LTD	<u>2.5MG</u>	<u>A202170 001</u>	Nov 28, 2011
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<u>AB</u>			<u>5MG</u>	<u>A202170 002</u>	Nov 28, 2011
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<u>AB</u>	!		<u>10MG</u>	<u>A202170 003</u>	Nov 28, 2011
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<u>AB</u>		VINTAGE PHARMS LLC	<u>2.5MG</u>	<u>A200815 001</u>	Oct 28, 2011
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<u>AB</u>			<u>5MG</u>	<u>A200815 002</u>	Oct 28, 2011
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<u>AB</u>			<u>10MG</u>	<u>A200815 003</u>	Oct 28, 2011
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<u>AB</u>		YILING PHARM LTD	<u>2.5MG</u>	<u>A210847 001</u>	Oct 26, 2018
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<u>AB</u>			<u>5MG</u>	<u>A210847 002</u>	Oct 26, 2018
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<u>AB</u>			<u>10MG</u>	<u>A210847 003</u>	Oct 26, 2018
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<u>AB</u>		YUNG SHIN PHARM	<u>2.5MG</u>	<u>A204800 001</u>	Apr 29, 2019
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<u>AB</u>			<u>5MG</u>	<u>A204800 002</u>	Apr 29, 2019
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PRESCRIPTION DRUG PRODUCT LIST

FELODIPINE

TABLET, EXTENDED RELEASE;ORAL

FELODIPINE

<u>AB</u>		<u>10MG</u>	<u>A204800 003</u>	Apr 29, 2019
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FENOFIBRATE

CAPSULE;ORAL

ANTARA (MICRONIZED)

<u>AB</u>	+	LUPIN ATLANTIS	<u>43MG</u>	<u>N021695 001</u>	Nov 30, 2004
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<u>AB</u>	+	!	<u>130MG</u>	<u>N021695 003</u>	Nov 30, 2004
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FENOFIBRATE

<u>AB</u>		SUN PHARM INDS LTD	<u>43MG</u>	<u>A201748 001</u>	Oct 31, 2014
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<u>AB</u>			<u>130MG</u>	<u>A201748 002</u>	Oct 31, 2014
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FENOFIBRATE (MICRONIZED)

<u>AB</u>		AJANTA PHARMA LTD	<u>67MG</u>	<u>A210705 001</u>	Sep 10, 2018
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<u>AB</u>			<u>134MG</u>	<u>A210705 002</u>	Sep 10, 2018
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<u>AB</u>			<u>200MG</u>	<u>A210705 003</u>	Sep 10, 2018
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<u>AB</u>		ANI PHARMS INC	<u>67MG</u>	<u>A209504 001</u>	Apr 30, 2018
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<u>AB</u>			<u>134MG</u>	<u>A209504 002</u>	Apr 30, 2018
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<u>AB</u>			<u>200MG</u>	<u>A209504 003</u>	Apr 30, 2018
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<u>AB</u>		APOTEX	<u>43MG</u>	<u>A202252 001</u>	Jul 26, 2013
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<u>AB</u>			<u>130MG</u>	<u>A202252 002</u>	Jul 26, 2013
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<u>AB</u>		AUSTARPHARMA	<u>67MG</u>	<u>A207805 001</u>	Nov 16, 2017
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<u>AB</u>			<u>134MG</u>	<u>A207805 002</u>	Nov 16, 2017
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<u>AB</u>			<u>200MG</u>	<u>A207805 003</u>	Nov 16, 2017
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<u>AB</u>		DR REDDYS LABS SA	<u>43MG</u>	<u>A090859 001</u>	Mar 01, 2012
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<u>AB</u>			<u>130MG</u>	<u>A090859 002</u>	Mar 01, 2012
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<u>AB</u>		GLENMARK PHARMS LTD	<u>67MG</u>	<u>A205566 001</u>	Apr 07, 2017
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<u>AB</u>			<u>134MG</u>	<u>A205566 002</u>	Apr 07, 2017
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<u>AB</u>			<u>200MG</u>	<u>A205566 003</u>	Apr 07, 2017
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<u>AB</u>		IMPAX LABS	<u>67MG</u>	<u>A075868 001</u>	Oct 27, 2003
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<u>AB</u>			<u>134MG</u>	<u>A075868 002</u>	Oct 27, 2003
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<u>AB</u>	!		<u>200MG</u>	<u>A075868 003</u>	Oct 27, 2003
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<u>AB</u>		INVAGEN PHARMS	<u>67MG</u>	<u>A207378 001</u>	Mar 28, 2017
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<u>AB</u>			<u>134MG</u>	<u>A207378 002</u>	Mar 28, 2017
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<u>AB</u>			<u>200MG</u>	<u>A207378 003</u>	Mar 28, 2017
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<u>AB</u>		MYLAN PHARMS INC	<u>43MG</u>	<u>A202579 001</u>	Jan 10, 2013
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<u>AB</u>			<u>67MG</u>	<u>A202676 001</u>	Oct 23, 2012
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<u>AB</u>			<u>130MG</u>	<u>A202579 002</u>	Jan 10, 2013
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<u>AB</u>			<u>134MG</u>	<u>A202676 002</u>	Oct 23, 2012
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<u>AB</u>			<u>200MG</u>	<u>A202676 003</u>	Oct 23, 2012
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<u>AB</u>		RHODES PHARMS	<u>67MG</u>	<u>A075753 001</u>	Sep 03, 2002
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<u>AB</u>			<u>134MG</u>	<u>A075753 002</u>	Apr 09, 2002
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<u>AB</u>			<u>200MG</u>	<u>A075753 003</u>	Apr 09, 2002
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<u>AB</u>		TORRENT	<u>67MG</u>	<u>A210782 001</u>	Jun 26, 2018
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<u>AB</u>			<u>134MG</u>	<u>A210782 002</u>	Jun 26, 2018
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<u>AB</u>			<u>200MG</u>	<u>A210782 003</u>	Jun 26, 2018
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ANTARA (MICRONIZED)

	+	LUPIN ATLANTIS	30MG	N021695 004	Oct 18, 2013
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	+		90MG	N021695 005	Oct 18, 2013
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LIPOFEN

	+	CIPHER PHARMS INC	50MG	N021612 001	Jan 11, 2006
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	+	!	150MG	N021612 003	Jan 11, 2006
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TABLET;ORAL

FENOFIBRATE

<u>AB</u>		AJANTA PHARMA LTD	<u>54MG</u>	<u>A210138 001</u>	Jul 23, 2018
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<u>AB</u>			<u>160MG</u>	<u>A210138 002</u>	Jul 23, 2018
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<u>AB</u>		ALEMBIC PHARMS LTD	<u>48MG</u>	<u>A210476 001</u>	Aug 09, 2019
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<u>AB</u>			<u>54MG</u>	<u>A213252 001</u>	Jan 17, 2020
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<u>AB</u>			<u>145MG</u>	<u>A210476 002</u>	Aug 09, 2019
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<u>AB</u>			<u>160MG</u>	<u>A213252 002</u>	Jan 17, 2020
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<u>AB</u>		AMNEAL PHARMS LLC	<u>48MG</u>	<u>A209951 001</u>	Feb 09, 2018
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<u>AB</u>			<u>54MG</u>	<u>A209950 001</u>	Mar 19, 2018
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<u>AB</u>			<u>145MG</u>	<u>A209951 002</u>	Feb 09, 2018
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<u>AB</u>			<u>160MG</u>	<u>A209950 002</u>	Mar 19, 2018
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<u>AB</u>		APPCO	<u>54MG</u>	<u>A210670 001</u>	Sep 06, 2019
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<u>AB</u>			<u>160MG</u>	<u>A210670 002</u>	Sep 06, 2019
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<u>AB</u>		AUROBINDO PHARMA LTD	<u>48MG</u>	<u>A205118 001</u>	May 05, 2016
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<u>AB</u>			<u>145MG</u>	<u>A205118 002</u>	May 05, 2016
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<u>AB</u>		AUSTARPHARMA	<u>54MG</u>	<u>A207803 001</u>	Dec 19, 2017
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<u>AB</u>			<u>160MG</u>	<u>A207803 002</u>	Dec 19, 2017
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<u>AB</u>		CIPLA	<u>48MG</u>	<u>A208709 001</u>	Dec 15, 2016
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PRESCRIPTION DRUG PRODUCT LISTFENOFIBRATE

TABLET; ORAL

FENOFIBRATE

<u>AB</u>		<u>145MG</u>	<u>A208709 002</u>	Dec 15, 2016
<u>AB</u>	GRAVITI PHARMS	<u>48MG</u>	<u>A211122 001</u>	Mar 18, 2020
<u>AB</u>		<u>54MG</u>	<u>A210606 001</u>	Aug 17, 2018
<u>AB</u>		<u>145MG</u>	<u>A211122 002</u>	Mar 18, 2020
<u>AB</u>		<u>160MG</u>	<u>A210606 002</u>	Aug 17, 2018
<u>AB</u>	HETERO LABS LTD III	<u>48MG</u>	<u>A204598 001</u>	Jul 12, 2016
<u>AB</u>		<u>145MG</u>	<u>A204598 002</u>	Jul 12, 2016
<u>AB</u>	IMPAX LABS	<u>54MG</u>	<u>A076509 001</u>	Mar 26, 2008
<u>AB</u>	!	<u>160MG</u>	<u>A076509 002</u>	Mar 26, 2008
<u>AB</u>	LUPIN LTD	<u>48MG</u>	<u>A090856 001</u>	Dec 23, 2011
<u>AB</u>		<u>54MG</u>	<u>A204019 001</u>	Aug 17, 2015
<u>AB</u>		<u>145MG</u>	<u>A090856 002</u>	Dec 23, 2011
<u>AB</u>		<u>160MG</u>	<u>A204019 002</u>	Aug 17, 2015
<u>AB</u>	MYLAN	<u>40MG</u>	<u>A204475 001</u>	Jun 23, 2016
<u>AB</u>		<u>54MG</u>	<u>A076520 001</u>	Oct 25, 2007
<u>AB</u>		<u>120MG</u>	<u>A204475 002</u>	Jun 23, 2016
<u>AB</u>		<u>160MG</u>	<u>A076520 003</u>	Oct 25, 2007
<u>AB</u>	MYLAN PHARMS INC	<u>48MG</u>	<u>A202856 001</u>	Dec 07, 2012
<u>AB</u>		<u>145MG</u>	<u>A202856 002</u>	Dec 07, 2012
<u>AB</u>	ORIT LABS LLC	<u>54MG</u>	<u>A209660 001</u>	Feb 11, 2019
<u>AB</u>		<u>160MG</u>	<u>A209660 002</u>	Feb 11, 2019
<u>AB</u>	PRINSTON INC	<u>48MG</u>	<u>A211080 001</u>	Aug 28, 2018
<u>AB</u>		<u>145MG</u>	<u>A211080 002</u>	Aug 28, 2018
<u>AB</u>	RHODES PHARMS	<u>54MG</u>	<u>A076433 001</u>	May 13, 2005
<u>AB</u>		<u>160MG</u>	<u>A076433 002</u>	May 13, 2005
<u>AB</u>	SUN PHARM	<u>48MG</u>	<u>A200884 001</u>	Sep 07, 2017
<u>AB</u>		<u>145MG</u>	<u>A200884 002</u>	Sep 07, 2017
<u>AB</u>	SUN PHARM INDS LTD	<u>54MG</u>	<u>A076635 001</u>	Oct 31, 2005
<u>AB</u>		<u>160MG</u>	<u>A076635 003</u>	Oct 31, 2005
<u>AB</u>	VALEANT PHARMS NORTH	<u>48MG</u>	<u>A090715 001</u>	Apr 05, 2012
<u>AB</u>		<u>145MG</u>	<u>A090715 002</u>	Apr 05, 2012
	<u>FENOGLIDE</u>			
<u>AB</u>	+ SALIX	<u>40MG</u>	<u>N022118 001</u>	Aug 10, 2007
<u>AB</u>	+!	<u>120MG</u>	<u>N022118 002</u>	Aug 10, 2007
	<u>TRICOR</u>			
<u>AB</u>	+ ABBVIE	<u>48MG</u>	<u>N021656 001</u>	Nov 05, 2004
<u>AB</u>	+!	<u>145MG</u>	<u>N021656 002</u>	Nov 05, 2004
	TRIGLIDE			
BX	+ SKYEPHARMA AG	160MG	N021350 002	May 07, 2005
	FENOFIBRATE			
	SUN PHARM INDS LTD	107MG	A076635 002	Oct 31, 2005

FENOFIBRIC ACID

TABLET; ORAL

FIBRICOR

+ ATHENA
+!

35MG
105MG

N022418 001 Aug 14, 2009
N022418 002 Aug 14, 2009

FENOLDOPAM MESYLATE

INJECTABLE; INJECTION

CORLOPAMAP +! HOSPIRAEQ 10MG BASE/MLN019922 001 Sep 23, 1997FENOLDOPAM MESYLATEAP SANDOZ INCEQ 10MG BASE/MLA077155 001 Feb 15, 2005AP WEST-WARD PHARMS
INTEQ 10MG BASE/MLA076582 001 Oct 12, 2004FENOPROFEN CALCIUM

CAPSULE; ORAL

NALFON

+ XSPIRE PHARMA
+!

EQ 200MG BASE
EQ 400MG BASE

N017604 003
N017604 004 Jul 21, 2009

TABLET; ORAL

FENOPROFEN CALCIUM

! XSPIRE PHARMA

EQ 600MG BASE

A072267 001 Aug 17, 1988

PRESCRIPTION DRUG PRODUCT LIST

FENTANYL

FILM, EXTENDED RELEASE; TRANSDERMAL

<u>DURAGESIC-100</u>						
AB	+	JANSSEN PHARMS	100MCG/HR	N019813	001	Aug 07, 1990
<u>DURAGESIC-12</u>						
AB	+	JANSSEN PHARMS	12.5MCG/HR	N019813	005	Feb 04, 2005
<u>DURAGESIC-25</u>						
AB	+	JANSSEN PHARMS	25MCG/HR	N019813	004	Aug 07, 1990
<u>DURAGESIC-37</u>						
AB	+	JANSSEN PHARMS	37.5MCG/HR	N019813	006	Jan 24, 2018
<u>DURAGESIC-50</u>						
AB	+	JANSSEN PHARMS	50MCG/HR	N019813	003	Aug 07, 1990
<u>DURAGESIC-75</u>						
AB	+	JANSSEN PHARMS	75MCG/HR	N019813	002	Aug 07, 1990
<u>FENTANYL-100</u>						
AB		3M DRUG DELIVERY	100MCG/HR	A202097	005	Nov 04, 2016
AB		AVEVA	100MCG/HR	A077449	004	Oct 20, 2008
AB		LAVIPHARM LABS	100MCG/HR	A077051	004	Aug 04, 2006
AB		MYLAN TECHNOLOGIES	100MCG/HR	A076258	004	Jan 28, 2005
AB		SPECGX LLC	100MCG/HR	A077154	004	Feb 09, 2011
<u>FENTANYL-12</u>						
AB		3M DRUG DELIVERY	12.5MCG/HR	A202097	001	Nov 04, 2016
AB		AVEVA	12.5MCG/HR	A077449	005	Sep 11, 2015
AB		MYLAN TECHNOLOGIES	12.5MCG/HR	A076258	005	Jan 23, 2007
AB		SPECGX LLC	12.5MCG/HR	A077154	005	Jun 11, 2015
<u>FENTANYL-25</u>						
AB		3M DRUG DELIVERY	25MCG/HR	A202097	002	Nov 04, 2016
AB		AVEVA	25MCG/HR	A077449	001	Oct 20, 2008
AB		LAVIPHARM LABS	25MCG/HR	A077051	001	Aug 04, 2006
AB		MYLAN TECHNOLOGIES	25MCG/HR	A076258	001	Jan 28, 2005
AB		SPECGX LLC	25MCG/HR	A077154	001	Feb 09, 2011
<u>FENTANYL-37</u>						
AB		AVEVA	37.5MCG/HR	A077449	006	Dec 06, 2017
AB		MYLAN TECHNOLOGIES	37.5MCG/HR	A076258	006	Dec 29, 2014
<u>FENTANYL-50</u>						
AB		3M DRUG DELIVERY	50MCG/HR	A202097	003	Nov 04, 2016
AB		AVEVA	50MCG/HR	A077449	002	Oct 20, 2008
AB		LAVIPHARM LABS	50MCG/HR	A077051	002	Aug 04, 2006
AB		MYLAN TECHNOLOGIES	50MCG/HR	A076258	002	Jan 28, 2005
AB		SPECGX LLC	50MCG/HR	A077154	002	Feb 09, 2011
<u>FENTANYL-62</u>						
AB		AVEVA	62.5MCG/HR	A077449	007	Dec 06, 2017
<u>FENTANYL-75</u>						
AB		3M DRUG DELIVERY	75MCG/HR	A202097	004	Nov 04, 2016
AB		AVEVA	75MCG/HR	A077449	003	Oct 20, 2008
AB		LAVIPHARM LABS	75MCG/HR	A077051	003	Aug 04, 2006
AB		MYLAN TECHNOLOGIES	75MCG/HR	A076258	003	Jan 28, 2005
AB		SPECGX LLC	75MCG/HR	A077154	003	Feb 09, 2011
<u>FENTANYL-87</u>						
AB		AVEVA	87.5MCG/HR	A077449	008	Dec 06, 2017
		FENTANYL-62				
		MYLAN TECHNOLOGIES	62.5MCG/HR	A076258	007	Dec 29, 2014
		FENTANYL-87				
		MYLAN TECHNOLOGIES	87.5MCG/HR	A076258	008	Dec 29, 2014
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
<u>FENTANYL CITRATE</u>						
INJECTABLE; INJECTION						
<u>FENTANYL CITRATE</u>						
AP		HOSPIRA	EQ 0.05MG BASE/ML	N019115	001	Jan 12, 1985
<u>FENTANYL CITRATE PRESERVATIVE FREE</u>						
AP		FRESENIUS KABI USA	EQ 0.05MG BASE/ML	A210762	001	May 03, 2019
AP	+	HIKMA	EQ 0.05MG BASE/ML	N019101	001	Jul 11, 1984
AP		HOSPIRA	EQ 0.05MG BASE/ML	A072786	001	Sep 24, 1991

PRESCRIPTION DRUG PRODUCT LIST

FENTANYL CITRATE

INJECTABLE; INJECTION

SUBLIMAZE PRESERVATIVE FREE

AP	+ !	AKORN	EQ 0.05MG BASE/ML	N016619 001	
		SPRAY, METERED; NASAL			
		LAZANDA			
	+	BTCP PHARMA	EQ 0.1MG BASE	N022569 001	Jun 30, 2011
	+ !		EQ 0.4MG BASE	N022569 002	Jun 30, 2011
		TABLET; BUCCAL, SUBLINGUAL			
		FENTORA			
	+	CEPHALON	EQ 0.1MG BASE	N021947 001	Sep 25, 2006
	+		EQ 0.2MG BASE	N021947 002	Sep 25, 2006
	+ !		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
		TROCHE/LOZENGE; TRANSMUCOSAL			

ACTIQ

AB	+	CEPHALON	EQ 0.2MG BASE	N020747 001	Nov 04, 1998
AB	+ !		EQ 0.4MG BASE	N020747 002	Nov 04, 1998
AB	+		EQ 0.6MG BASE	N020747 003	Nov 04, 1998
AB	+		EQ 0.8MG BASE	N020747 004	Nov 04, 1998
AB	+		EQ 1.2MG BASE	N020747 005	Nov 04, 1998
AB	+		EQ 1.6MG BASE	N020747 006	Nov 04, 1998

FENTANYL CITRATE

AB		SPECGX LLC	EQ 0.2MG BASE	A078907 001	Oct 30, 2009
AB			EQ 0.4MG BASE	A078907 002	Oct 30, 2009
AB			EQ 0.6MG BASE	A078907 003	Oct 30, 2009
AB			EQ 0.8MG BASE	A078907 004	Oct 30, 2009
AB			EQ 1.2MG BASE	A078907 005	Oct 30, 2009
AB			EQ 1.6MG BASE	A078907 006	Oct 30, 2009

FERRIC CARBOXYMALTOSE

INJECTABLE; INTRAVENOUS

INJECTAFER

+ !	AM REGENT	750MG IRON/15ML (50MG IRON/ML)	N203565 001	Jul 25, 2013
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FERRIC CITRATE

TABLET; ORAL

AURYXIA

+ !	KERYX BIOPHARMS	EQ 210MG IRON	N205874 001	Sep 05, 2014
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FERRIC DERISOMALTOSE

SOLUTION; INTRAVENOUS

MONOFERRIC

+ !	PHARMACOSMOS AS	100MG/ML (100MG/ML)	N208171 001	Jan 16, 2020
+ !		500MG/5ML (100MG/ML)	N208171 002	Jan 16, 2020
+ !		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
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FERRIC MALTOI

CAPSULE; ORAL

ACCRUFER

+ !	SHIELD TX	30MG IRON	N212320 001	Jul 25, 2019
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FERRIC PYROPHOSPHATE CITRATE

FOR SOLUTION; INTRAVENOUS

TRIFERIC

+ !	ROCKWELL MEDICAL INC	272MG IRON/PACKET	N208551 001	Apr 25, 2016
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SOLUTION; INTRAVENOUS

TRIFERIC

+ !	ROCKWELL MEDICAL INC	27.2MG IRON/5ML (5.44MG IRON/ML)	N206317 001	Jan 23, 2015
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FERUMOXYTOL

SOLUTION; INTRAVENOUS

FERAHEME

+ !	AMAG PHARMS INC	EQ 510MG IRON/17ML (EQ 30MG IRON/ML)	N022180 001	Jun 30, 2009
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PRESCRIPTION DRUG PRODUCT LIST

FESOTERODINE FUMARATE

TABLET, EXTENDED RELEASE;ORAL

FESOTERODINE FUMARATE

<u>AB</u>	AUROBINDO PHARMA LTD	<u>4MG</u>	<u>A205007 001</u>	Feb 17, 2017
<u>AB</u>		<u>8MG</u>	<u>A205007 002</u>	Feb 17, 2017
<u>AB</u>	ZYDUS PHARMS	<u>4MG</u>	<u>A204946 001</u>	Oct 03, 2017
<u>AB</u>		<u>8MG</u>	<u>A204946 002</u>	Oct 03, 2017
<u>TOVIAZ</u>				
<u>AB</u>	+ PFIZER	<u>4MG</u>	<u>N022030 001</u>	Oct 31, 2008
<u>AB</u>	+!	<u>8MG</u>	<u>N022030 002</u>	Oct 31, 2008

FEXOFENADINE HYDROCHLORIDE

TABLET;ORAL

FEXOFENADINE HYDROCHLORIDE

<u>AB</u>	BARR	<u>30MG</u>	<u>A076191 001</u>	Aug 31, 2005
<u>AB</u>		<u>60MG</u>	<u>A076191 002</u>	Aug 31, 2005
<u>AB</u>		<u>180MG</u>	<u>A076191 003</u>	Aug 31, 2005
<u>AB</u>	DR REDDYS LABS LTD	<u>30MG</u>	<u>A076502 001</u>	Apr 11, 2006
<u>AB</u>		<u>60MG</u>	<u>A076502 002</u>	Apr 11, 2006
<u>AB</u>		<u>180MG</u>	<u>A076502 003</u>	Apr 11, 2006
<u>AB</u>	MYLAN	<u>60MG</u>	<u>A077081 003</u>	Apr 11, 2008
<u>AB</u>		<u>180MG</u>	<u>A077081 001</u>	Apr 16, 2007
<u>AB</u>	TEVA	<u>30MG</u>	<u>A076447 001</u>	Sep 01, 2005
<u>AB</u>		<u>60MG</u>	<u>A076447 002</u>	Sep 01, 2005
<u>AB</u>		<u>180MG</u>	<u>A076447 003</u>	Sep 01, 2005

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

FINASTERIDE

<u>AB</u>	ACCORD HLTHCARE	<u>1MG</u>	<u>A091643 001</u>	Nov 05, 2013
<u>AB</u>		<u>5MG</u>	<u>A090121 001</u>	Feb 23, 2010
<u>AB</u>	ACTAVIS TOTOWA	<u>1MG</u>	<u>A078371 001</u>	Nov 05, 2013
<u>AB</u>	ACTAVIS TOTOWA TEVA	<u>5MG</u>	<u>A077914 001</u>	Mar 28, 2007
<u>AB</u>	ALKEM LABS LTD	<u>1MG</u>	<u>A207750 001</u>	Jan 06, 2017
<u>AB</u>		<u>5MG</u>	<u>A204304 001</u>	Jan 05, 2017
<u>AB</u>	AUROBINDO PHARMA	<u>5MG</u>	<u>A078341 001</u>	Oct 30, 2007
<u>AB</u>	AUROBINDO PHARMA LTD	<u>1MG</u>	<u>A203687 001</u>	Nov 05, 2013
<u>AB</u>	CIPLA	<u>1MG</u>	<u>A077335 001</u>	Nov 20, 2014
<u>AB</u>	DR REDDYS LABS INC	<u>1MG</u>	<u>A076436 001</u>	Jul 28, 2006
<u>AB</u>	DR REDDYS LABS LTD	<u>5MG</u>	<u>A076437 001</u>	Feb 28, 2007
<u>AB</u>	HETERO LABS LTD III	<u>1MG</u>	<u>A090060 001</u>	Jul 01, 2013
<u>AB</u>		<u>5MG</u>	<u>A090061 001</u>	Jun 07, 2010
<u>AB</u>	SUN PHARM	<u>1MG</u>	<u>A090508 001</u>	Jul 01, 2013
<u>AB</u>		<u>5MG</u>	<u>A090507 001</u>	Aug 16, 2011
<u>AB</u>	TEVA	<u>1MG</u>	<u>A076905 001</u>	Nov 05, 2013
<u>AB</u>		<u>5MG</u>	<u>A076511 001</u>	Dec 15, 2006
<u>AB</u>	ZYDUS PHARMS USA INC	<u>5MG</u>	<u>A078900 001</u>	Dec 28, 2009

PROPECIA

<u>AB</u>	+! MERCK	<u>1MG</u>	<u>N020788 001</u>	Dec 19, 1997
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PROSCAR

<u>AB</u>	+! MERCK	<u>5MG</u>	<u>N020180 001</u>	Jun 19, 1992
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FINGOLIMOD HYDROCHLORIDE

CAPSULE;ORAL

FINGOLIMOD HYDROCHLORIDE

<u>AB</u>	BIOCON LTD	<u>EQ 0.5MG BASE</u>	<u>A207979 001</u>	Dec 04, 2019
<u>AB</u>	HEC PHARM CO LTD	<u>EQ 0.5MG BASE</u>	<u>A207939 001</u>	Dec 04, 2019
<u>AB</u>	SUN PHARM	<u>EQ 0.5MG BASE</u>	<u>A208014 001</u>	Dec 04, 2019

GILENYA

<u>AB</u>	+! NOVARTIS	<u>EQ 0.5MG BASE</u>	<u>N022527 001</u>	Sep 21, 2010
	+	EQ 0.25MG BASE	N022527 002	May 11, 2018

PRESCRIPTION DRUG PRODUCT LISTFISH 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	! PADDOCK LLC	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 INC	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	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
TAMBOCOR					
AB	+ CNTY LINE PHARMS	50MG	N018830	004	Aug 23, 1988
AB	+	100MG	N018830	001	Oct 31, 1985
AB	+	150MG	N018830	003	Jun 03, 1988

FLIBANSERIN

TABLET; ORAL

ADDYI

+!	SPROUT PHARMS	100MG	N022526	001	Aug 18, 2015
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FLORBETABEN F-18

SOLUTION; INTRAVENOUS

NEURACEQ

+!	LIFE MOLECULAR	30ML (1.4-135mCi/ML)	N204677	001	Mar 19, 2014
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FLORBETAPIR F-18

SOLUTION; INTRAVENOUS

AMYVID

+!	AVID RADIOPHARMS INC	10-30ML (13.5-51mCi/ML)	N202008	002	Apr 06, 2012
+!		10-50ML (13.5-51mCi/ML)	N202008	003	Apr 06, 2012

FLOXURIDINE

INJECTABLE; INJECTION

FLOXURIDINE

AP	FRESENIUS KABI USA	500MG/VIAL	A075837	001	Feb 22, 2001
AP	! WEST-WARD PHARMS INT	500MG/VIAL	A075387	001	Apr 16, 2000

PRESCRIPTION DRUG PRODUCT LIST

FLUCICLOVINE 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

LTD

AB 200MG/5ML **A079150 002** Sep 18, 2009

INJECTABLE; INJECTION

FLUCONAZOLE IN DEXTROSE 5% IN PLASTIC CONTAINER

AP HIKMA FARMACEUTICA 200MG/100ML (2MG/ML) **A078764 001** Jan 30, 2012

AP 400MG/200ML (2MG/ML) **A078764 002** Jan 30, 2012

AP ! HOSPIRA 200MG/100ML (2MG/ML) **A076304 001** Jul 29, 2004

AP ! 400MG/200ML (2MG/ML) **A076304 002** Jul 29, 2004

AP WOODWARD 200MG/100ML (2MG/ML) **A077988 001** May 26, 2010

AP 400MG/200ML (2MG/ML) **A077988 002** May 26, 2010

FLUCONAZOLE IN SODIUM CHLORIDE 0.9%

AP BAXTER HLTHCARE 200MG/100ML (2MG/ML) **A077947 001** May 26, 2010

CORP

AP 400MG/200ML (2MG/ML) **A077947 002** May 26, 2010

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 FARMACEUTICA 200MG/100ML (2MG/ML) **A076736 001** Aug 23, 2005

AP WEST-WARD PHARMS 200MG/100ML (2MG/ML) **A076087 001** Jul 29, 2004

INT

AP 400MG/200ML (2MG/ML) **A076087 003** Jul 29, 2004

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 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 WEST-WARD PHARMS 200MG/100ML (2MG/ML) **A078107 001** Jul 30, 2008

INT

AP 400MG/200ML (2MG/ML) **A078107 002** Jul 30, 2008

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%

WEST-WARD PHARMS 100MG/50ML (2MG/ML)

A076087 002 Sep 26, 2008

INT

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

FLUCONAZOLE

AB AUROBINDO PHARMA 50MG **A077731 001** Oct 07, 2008

AB 100MG **A077731 002** Oct 07, 2008

AB 150MG **A077731 003** Oct 07, 2008

AB 200MG **A077731 004** Oct 07, 2008

AB CHARTWELL 50MG **A076665 001** Jul 29, 2004

AB 100MG **A076665 002** Jul 29, 2004

AB 150MG **A076665 003** Jul 29, 2004

AB 200MG **A076665 004** Jul 29, 2004

AB DR REDDYS LABS INC 50MG **A076658 001** Jul 29, 2004

AB 100MG **A076658 002** Jul 29, 2004

AB 150MG **A076658 003** Jul 29, 2004

AB 200MG **A076658 004** Jul 29, 2004

AB GLENMARK GENERICS 50MG **A077253 001** Jan 25, 2006

AB 100MG **A077253 002** Jan 25, 2006

AB 150MG **A077253 003** Jan 25, 2006

AB 200MG **A077253 004** Jan 25, 2006

PRESCRIPTION DRUG PRODUCT LIST

3-191 (of 453)

FLUCONAZOLE

TABLET; ORAL

FLUCONAZOLE

<u>AB</u>	HARRIS PHARM	<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>	IVAX SUB TEVA PHARMS	<u>50MG</u>	<u>A076077 001</u>	Jul 29, 2004
<u>AB</u>		<u>100MG</u>	<u>A076077 002</u>	Jul 29, 2004
<u>AB</u>		<u>150MG</u>	<u>A076077 003</u>	Jul 29, 2004
<u>AB</u>		<u>200MG</u>	<u>A076077 004</u>	Jul 29, 2004
<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>	TEVA	<u>50MG</u>	<u>A074681 001</u>	Jul 29, 2004
<u>AB</u>		<u>100MG</u>	<u>A074681 002</u>	Jul 29, 2004
<u>AB</u>		<u>150MG</u>	<u>A074681 003</u>	Jul 29, 2004
<u>AB</u>		<u>200MG</u>	<u>A074681 004</u>	Jul 29, 2004
<u>AB</u>	UNIQUE PHARM LABS	<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

FLUCYTOSINE

CAPSULE; ORAL

ANCOBON

<u>AB</u>	+ BAUSCH	<u>250MG</u>	<u>N017001 001</u>	
<u>AB</u>	+!	<u>500MG</u>	<u>N017001 002</u>	

FLUCYTOSINE

<u>AB</u>	HIKMA	<u>250MG</u>	<u>A206550 001</u>	Oct 17, 2017
<u>AB</u>		<u>500MG</u>	<u>A206550 002</u>	Oct 17, 2017
<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>	RECIPHARM	<u>250MG</u>	<u>A207536 001</u>	Jun 18, 2018
<u>AB</u>		<u>500MG</u>	<u>A207536 002</u>	Jun 18, 2018
<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

FLUDARABINE PHOSPHATE

INJECTABLE; INJECTION

FLUDARABINE PHOSPHATE

<u>AP</u>	ACTAVIS LLC	<u>50MG/2ML (25MG/ML)</u>	<u>A203738 001</u>	Feb 28, 2017
<u>AP</u>	ACTAVIS TOTOWA	<u>50MG/VIAL</u>	<u>A078610 001</u>	Feb 11, 2009
<u>AP</u>	AREVA PHARMS	<u>50MG/2ML (25MG/ML)</u>	<u>A090724 001</u>	Sep 27, 2010
<u>AP</u>	CUSTOPHARM INC	<u>50MG/VIAL</u>	<u>A076349 001</u>	Aug 28, 2003
<u>AP</u>	! FRESENIUS KABI USA	<u>50MG/2ML (25MG/ML)</u>	<u>A078393 001</u>	Oct 15, 2007
<u>AP</u>		<u>50MG/VIAL</u>	<u>A078544 001</u>	Oct 15, 2007
<u>AP</u>	! HOSPIRA	<u>50MG/VIAL</u>	<u>A077790 001</u>	Apr 06, 2007
<u>AP</u>	MYLAN LABS LTD	<u>50MG/VIAL</u>	<u>A200648 001</u>	Oct 16, 2012
<u>AP</u>	SAGENT PHARMS INC	<u>50MG/2ML (25MG/ML)</u>	<u>A076661 001</u>	Apr 28, 2004
<u>AP</u>	+! SANDOZ	<u>50MG/2ML (25MG/ML)</u>	<u>N022137 001</u>	Sep 21, 2007

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>	CHILDRENS HOSP MI	<u>20-300mCi/ML</u>	<u>A204385 001</u>	Oct 29, 2014
<u>AP</u>	CPDC	<u>20-300mCi/ML</u>	<u>A204525 001</u>	Oct 29, 2014
<u>AP</u>	DECATUR	<u>20-300mCi/ML</u>	<u>A204463 001</u>	Oct 21, 2014
<u>AP</u>	ESSENTIAL ISOTOPES	<u>20-300mCi/ML</u>	<u>A203946 001</u>	Feb 05, 2014
<u>AP</u>	+! FEINSTEIN	<u>20-400mCi/ML</u>	<u>N021870 002</u>	Nov 21, 2008
<u>AP</u>	JUBILANT DRAXIMAGE	<u>20-300mCi/ML</u>	<u>A203920 001</u>	Jun 23, 2015
<u>AP</u>	KETTERING MEDCTR	<u>4-40mCi/ML</u>	<u>A204759 001</u>	Oct 27, 2015
<u>AP</u>	KREITCHMAN PET CTR	<u>10-100mCi/ML</u>	<u>A203942 001</u>	Apr 11, 2016

PRESCRIPTION DRUG PRODUCT LIST

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FLUDEOXYGLUCOSE F-18

INJECTABLE; INTRAVENOUS

FLUDEOXYGLUCOSE F18

<u>AP</u>	LANTHEUS MEDICAL	<u>20-200mCi/ML</u>	<u>A203664</u>	<u>001</u>	Feb 04, 2014
<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>	! QUEEN HAMAMATSU PET	<u>10-100mCi/ML</u>	<u>A203771</u>	<u>001</u>	Aug 31, 2015
<u>AP</u>	SHERTECH LABS LLC	<u>20-300mCi/ML</u>	<u>A204264</u>	<u>001</u>	Dec 18, 2014
<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>20-500mCi/ML</u>	<u>A203665</u>	<u>001</u>	Feb 14, 2013
<u>AP</u>	! UCLA BIOMEDICAL	<u>20-200mCi/ML</u>	<u>A203801</u>	<u>001</u>	Oct 29, 2014
<u>AP</u>	! UCSF RODIOPHARM	<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 TX MD ANDERSON	<u>20-300mCi/ML</u>	<u>A203246</u>	<u>002</u>	Jan 13, 2014
<u>AP</u>	UNIV TX SW MEDCTR	<u>20-200mCi/ML</u>	<u>A210265</u>	<u>001</u>	Feb 06, 2020
<u>AP</u>	UNIV UTAH CYCLOTRON	<u>20-300mCi/ML</u>	<u>A204498</u>	<u>001</u>	Jun 23, 2015
<u>AP</u>	WI MEDCL CYCLOTRON	<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
	HOT SHOTS NM LLC	4-500mCi/ML	A203937	001	Oct 30, 2014
	NORTHLAND	4-500mCi/ML	A203994	001	Feb 04, 2015
	PRECISION NUCLEAR	20-500mCi/ML	A204546	001	Apr 07, 2015
	SPECTRON MRC LLC	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>	HIKMA PHARMS	<u>0.1MG</u>	<u>A091302</u>	<u>001</u>	Jul 22, 2011
<u>AB</u>	! IMPAX LABS	<u>0.1MG</u>	<u>A040431</u>	<u>001</u>	Mar 18, 2002

FLUMAZENIL

INJECTABLE; INJECTION

FLUMAZENIL

<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 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>	MYLAN LABS LTD	<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 INC	<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
<u>AP</u>	WEST-WARD PHARMS	<u>0.5MG/5ML (0.1MG/ML)</u>	<u>A076256</u>	<u>002</u>	Oct 12, 2004
	INT				
<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

FLUNISOLIDE

SPRAY, METERED; NASAL

FLUNISOLIDE

<u>AB</u>	! BAUSCH AND LOMB	<u>0.025MG/SPRAY</u>	<u>A074805</u>	<u>001</u>	Feb 20, 2002
<u>AB</u>	HI TECH PHARMA CO	<u>0.025MG/SPRAY</u>	<u>A077704</u>	<u>001</u>	Aug 03, 2006

FLUOCINOLONE ACETONIDE

CREAM; TOPICAL

FLUOCINOLONE ACETONIDE

<u>AT</u>	ACP NIMBLE	<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
<u>AT</u>	TARO	<u>0.025%</u>	<u>A087104</u>	<u>001</u>	Apr 27, 1982
	<u>SYNALAR</u>				
<u>AT</u>	+! MEDIMETRIKS PHARMS	<u>0.01%</u>	<u>N012787</u>	<u>004</u>	
<u>AT</u>	+!	<u>0.025%</u>	<u>N012787</u>	<u>002</u>	
<u>AT</u>	+!	<u>0.025%</u>	<u>N012787</u>	<u>005</u>	

PRESCRIPTION DRUG PRODUCT LIST

FLUOCINOLONE ACETONIDE

IMPLANT; INTRAVITREAL

ILUVIEN

+	ALIMERA SCIENCES INC	0.19MG	N201923	001	Sep 26, 2014
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RETISERT

+	BAUSCH AND LOMB	0.59MG	N021737	001	Apr 08, 2005
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YUTIQ

+	EYEPOINT PHARMS	0.18MG	N210331	001	Oct 12, 2018
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OIL; TOPICAL

DERMA-SMOOTH/FS

<u>AT</u>	+	HILL DERMAC	<u>0.01%</u>	<u>N019452</u>	<u>001</u>	Feb 03, 1988
<u>AT</u>	+		<u>0.01%</u>	<u>N019452</u>	<u>002</u>	Nov 09, 2005

FLUCINOLONE ACETONIDE

<u>AT</u>		GLENMARK PHARMS LTD	<u>0.01%</u>	<u>A210556</u>	<u>001</u>	Oct 25, 2018
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FLUOCINOLONE ACETONIDE

<u>AT</u>		AKORN	<u>0.01%</u>	<u>A091514</u>	<u>001</u>	Jun 25, 2015
<u>AT</u>		IDENTI PHARMS INC	<u>0.01%</u>	<u>A201759</u>	<u>001</u>	Oct 17, 2011
<u>AT</u>			<u>0.01%</u>	<u>A201764</u>	<u>001</u>	Oct 17, 2011
<u>AT</u>		LYNE	<u>0.01%</u>	<u>A090982</u>	<u>001</u>	Apr 25, 2016
<u>AT</u>			<u>0.01%</u>	<u>A203377</u>	<u>001</u>	Apr 25, 2016
<u>AT</u>		PERRIGO ISRAEL	<u>0.01%</u>	<u>A202847</u>	<u>001</u>	Aug 09, 2013
<u>AT</u>			<u>0.01%</u>	<u>A202848</u>	<u>001</u>	Aug 09, 2013
<u>AT</u>		TARO	<u>0.01%</u>	<u>A202368</u>	<u>001</u>	May 19, 2016
<u>AT</u>			<u>0.01%</u>	<u>A209336</u>	<u>001</u>	May 19, 2016

FLUOCINONIDE ACETONIDE

<u>AT</u>		GLENMARK PHARMS LTD	<u>0.01%</u>	<u>A210539</u>	<u>001</u>	Oct 26, 2018
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OIL/DROPS; OTIC

DERMOTIC

<u>AT</u>	+	HILL DERMAC	<u>0.01%</u>	<u>N019452</u>	<u>003</u>	Nov 09, 2005
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FLAC

<u>AT</u>		ANDA REPOSITORY	<u>0.01%</u>	<u>A210736</u>	<u>001</u>	Apr 11, 2018
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FLUOCINOLONE ACETONIDE

<u>AT</u>		AKORN	<u>0.01%</u>	<u>A202705</u>	<u>001</u>	Sep 09, 2016
<u>AT</u>		IDENTI PHARMS INC	<u>0.01%</u>	<u>A091306</u>	<u>001</u>	Oct 17, 2011
<u>AT</u>		LYNE	<u>0.01%</u>	<u>A203378</u>	<u>001</u>	Apr 25, 2016
<u>AT</u>		PERRIGO ISRAEL	<u>0.01%</u>	<u>A202849</u>	<u>001</u>	Jul 17, 2017

FLUOCINONIDE ACETONIDE

<u>AT</u>		GLENMARK PHARMS LTD	<u>0.01%</u>	<u>A211815</u>	<u>001</u>	Dec 14, 2018
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OINTMENT; TOPICAL

FLUOCINOLONE ACETONIDE

<u>AT</u>		ACP NIMBLE	<u>0.025%</u>	<u>A089524</u>	<u>001</u>	Jul 26, 1988
<u>AT</u>		FOUGERA PHARMS INC	<u>0.025%</u>	<u>A088168</u>	<u>001</u>	Dec 16, 1982
<u>AT</u>		TARO	<u>0.025%</u>	<u>A040041</u>	<u>001</u>	Sep 15, 1994

SYNALAR

<u>AT</u>	+	MEDIMETRIKS PHARMS	<u>0.025%</u>	<u>N013960</u>	<u>001</u>	
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SHAMPOO; TOPICAL

CAPEX

+	GALDERMA LABS LP	0.01%	N020001	001	Aug 27, 1990
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SOLUTION; TOPICAL

FLUOCINOLONE ACETONIDE

<u>AT</u>		ENCUBE ETHICALS	<u>0.01%</u>	<u>A209913</u>	<u>001</u>	Feb 13, 2019
<u>AT</u>		FOUGERA PHARMS INC	<u>0.01%</u>	<u>A088167</u>	<u>001</u>	Dec 16, 1982
<u>AT</u>		GLASSHOUSE PHARMS	<u>0.01%</u>	<u>A209596</u>	<u>001</u>	Dec 26, 2017
<u>AT</u>		LUPIN	<u>0.01%</u>	<u>A206422</u>	<u>001</u>	Sep 02, 2015
<u>AT</u>		TARO	<u>0.01%</u>	<u>A089124</u>	<u>001</u>	Sep 11, 1985

SYNALAR

<u>AT</u>	+	MEDIMETRIKS PHARMS	<u>0.01%</u>	<u>N015296</u>	<u>001</u>	
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FLUOCINOLONE ACETONIDE; HYDROQUINONE; TRETINOIN

CREAM; TOPICAL

TRI-LUMA

+	GALDERMA LABS LP	0.01%; 4%; 0.05%	N021112	001	Jan 18, 2002
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FLUOCINOLONE ACETONIDE; NEOMYCIN SULFATE

CREAM; TOPICAL

NEO-SYNALAR

!	MEDIMETRIKS PHARMS	0.025%; EQ 3.5MG BASE/GM	A060700	001	
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PRESCRIPTION DRUG PRODUCT LIST

FLUOCINONIDE

CREAM; TOPICAL

FLUOCINONIDE

<u>AB</u>	AMNEAL PHARMS LLC	<u>0.1%</u>	<u>A211111</u>	<u>001</u>	Jun 04, 2018
<u>AB</u>	CADILA	<u>0.1%</u>	<u>A208989</u>	<u>001</u>	Feb 10, 2020
<u>AB</u>	FOUGERA PHARMS INC	<u>0.1%</u>	<u>A200735</u>	<u>001</u>	Jul 14, 2014
<u>AB</u>	GLENMARK GENERICS	<u>0.1%</u>	<u>A091282</u>	<u>001</u>	Jul 14, 2014
<u>AB</u>	PERRIGO ISRAEL	<u>0.1%</u>	<u>A090256</u>	<u>001</u>	Jan 14, 2014
<u>AB</u>	TARO	<u>0.1%</u>	<u>A200734</u>	<u>001</u>	Jul 14, 2014
<u>AB</u>	TELIGENT PHARMA INC	<u>0.1%</u>	<u>A211758</u>	<u>001</u>	Apr 03, 2019

VANOS

<u>AB</u>	+! MEDICIS	<u>0.1%</u>	<u>N021758</u>	<u>001</u>	Feb 11, 2005
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FLUOCINONIDE

<u>AB1</u>	ACP NIMBLE	<u>0.05%</u>	<u>A073085</u>	<u>001</u>	Feb 14, 1992
<u>AB1</u>	FOUGERA PHARMS INC	<u>0.05%</u>	<u>A073030</u>	<u>001</u>	Oct 17, 1994
<u>AB1</u>	TARO	<u>0.05%</u>	<u>A071500</u>	<u>001</u>	Jun 10, 1987
<u>AB1</u>	+!	<u>0.05%</u>	<u>N019117</u>	<u>001</u>	Jun 26, 1984
<u>AB1</u>	TELIGENT PHARMA INC	<u>0.05%</u>	<u>A211410</u>	<u>001</u>	Oct 16, 2018
<u>AB1</u>	TEVA	<u>0.05%</u>	<u>A072488</u>	<u>001</u>	Feb 06, 1989

FLUOCINONIDE EMULSIFIED BASE

<u>AB2</u>	ACP NIMBLE	<u>0.05%</u>	<u>A074204</u>	<u>001</u>	Jun 13, 1995
<u>AB2</u>	FOUGERA PHARMS	<u>0.05%</u>	<u>A076586</u>	<u>001</u>	Jun 23, 2004
<u>AB2</u>	! TARO PHARM INDS LTD	<u>0.05%</u>	<u>A072494</u>	<u>001</u>	Jan 19, 1989
<u>AB2</u>	TEVA	<u>0.05%</u>	<u>A072490</u>	<u>001</u>	Feb 07, 1989

LIDEX-E

<u>AB2</u>	+ CNTY LINE PHARMS	<u>0.05%</u>	<u>N016908</u>	<u>003</u>	
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GEL; TOPICAL

FLUOCINONIDE

<u>AB</u>	ACP NIMBLE	<u>0.05%</u>	<u>A072537</u>	<u>001</u>	Feb 07, 1989
<u>AB</u>	+ CNTY LINE PHARMS	<u>0.05%</u>	<u>N017373</u>	<u>001</u>	
<u>AB</u>	FOUGERA PHARMS INC	<u>0.05%</u>	<u>A072933</u>	<u>001</u>	Dec 30, 1994
<u>AB</u>	! TARO	<u>0.05%</u>	<u>A074935</u>	<u>001</u>	Jul 29, 1997
<u>AB</u>	TELIGENT PHARMA INC	<u>0.05%</u>	<u>A209030</u>	<u>001</u>	Jun 19, 2018

OINTMENT; TOPICAL

FLUOCINONIDE

<u>AB</u>	FOUGERA PHARMS	<u>0.05%</u>	<u>A074905</u>	<u>001</u>	Aug 26, 1997
<u>AB</u>	NOVEL LABS INC	<u>0.05%</u>	<u>A207538</u>	<u>001</u>	Jul 31, 2017
<u>AB</u>	! TARO	<u>0.05%</u>	<u>A075008</u>	<u>001</u>	Jun 30, 1999
<u>AB</u>	TELIGENT PHARMA INC	<u>0.05%</u>	<u>A207680</u>	<u>001</u>	Sep 28, 2018
<u>AB</u>	TEVA	<u>0.05%</u>	<u>A073481</u>	<u>001</u>	Dec 27, 1991
<u>AB</u>	XIROMED	<u>0.05%</u>	<u>A212976</u>	<u>001</u>	Nov 26, 2019

LIDEX

<u>AB</u>	+ CNTY LINE PHARMS	<u>0.05%</u>	<u>N016909</u>	<u>002</u>	
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SOLUTION; TOPICAL

FLUOCINONIDE

<u>AT</u>	ACP NIMBLE	<u>0.05%</u>	<u>A071535</u>	<u>001</u>	Dec 02, 1988
<u>AT</u>	ENCUBE ETHICALS	<u>0.05%</u>	<u>A209699</u>	<u>001</u>	Nov 29, 2018
<u>AT</u>	FOUGERA PHARMS INC	<u>0.05%</u>	<u>A072934</u>	<u>001</u>	Feb 27, 1995
<u>AT</u>	GLASSHOUSE PHARMS	<u>0.05%</u>	<u>A209118</u>	<u>001</u>	Apr 23, 2018
<u>AT</u>	MACLEODS PHARMS LTD	<u>0.05%</u>	<u>A209283</u>	<u>001</u>	Apr 23, 2018
<u>AT</u>	NOVEL LABS INC	<u>0.05%</u>	<u>A206003</u>	<u>001</u>	Jul 21, 2017
<u>AT</u>	! TARO	<u>0.05%</u>	<u>A074799</u>	<u>001</u>	Dec 31, 1996
<u>AT</u>	TELIGENT PHARMA INC	<u>0.05%</u>	<u>A207554</u>	<u>001</u>	Mar 18, 2019
<u>AT</u>	TEVA	<u>0.05%</u>	<u>A072511</u>	<u>001</u>	Feb 07, 1989
<u>AT</u>	ZYDUS PHARMS	<u>0.05%</u>	<u>A208948</u>	<u>001</u>	Jul 17, 2018

LIDEX

<u>AT</u>	+ CNTY LINE PHARMS	<u>0.05%</u>	<u>N018849</u>	<u>001</u>	Apr 06, 1984
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FLUORESCEIN SODIUM

INJECTABLE; INTRAVENOUS

AK-FLUOR 10%

<u>AP</u>	+ AKORN	<u>EQ 500MG BASE/5ML (EQ 100MG BASE/ML)</u>	<u>N022186</u>	<u>001</u>	Aug 08, 2008
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FLUORESCITE

<u>AP</u>	+! ALCON LABS INC	<u>EQ 500MG BASE/5ML (EQ 100MG BASE/ML)</u>	<u>N021980</u>	<u>001</u>	Mar 28, 2006
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AK-FLUOR 25%

	+! AKORN	<u>EQ 500MG BASE/2ML (EQ 250MG BASE/ML)</u>	<u>N022186</u>	<u>002</u>	Aug 08, 2008
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PRESCRIPTION DRUG PRODUCT LIST

FLUORODOPA F-18

SOLUTION; INTRAVENOUS

FLUORODOPA F18

+! FEINSTEIN 0.42-8.33mCi/ML N200655 001 Oct 10, 2019

FLUOROMETHOLONE

OINTMENT; OPHTHALMIC

FML

+! ALLERGAN 0.1% N017760 001 Sep 04, 1985

SUSPENSION/DROPS; OPHTHALMIC

FML

+! ALLERGAN 0.1% N016851 002 Jul 28, 1982

FML FORTE

+! ALLERGAN 0.25% N019216 001 Apr 23, 1986

FLUOROMETHOLONE ACETATE

SUSPENSION/DROPS; OPHTHALMIC

FLAREX

+! EYEVANCE 0.1% N019079 001 Feb 11, 1986

FLUOROURACIL

CREAM; TOPICAL

CARAC

AB +! VALEANT PHARMS NORTH **0.5%** **N020985 001** Oct 27, 2000

EFUDEX

AB +! BAUSCH **5%** **N016831 003**

FLUOROURACIL

AB MAYNE PHARMA **5%** **A077524 001** Apr 11, 2008

AB MYLAN **0.5%** **A203122 001** Apr 20, 2015

AB TARO **5%** **A090368 001** Mar 05, 2010

FLUOROPLEX

+! ALMIRALL 1% N016988 001

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 ! 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 INGENUS PHARMS LLC **500MG/10ML (50MG/ML)** **A209219 001** Dec 12, 2019

AP **500MG/10ML (50MG/ML)** **A209271 001** Dec 11, 2019

AP **1GM/20ML (50MG/ML)** **A209219 002** Dec 12, 2019

AP **2.5GM/50ML (50MG/ML)** **A209271 002** Dec 11, 2019

AP MYLAN LABS LTD **2.5GM/50ML (50MG/ML)** **A202669 001** Jul 17, 2012

AP **5GM/100ML (50MG/ML)** **A202669 002** Jul 17, 2012

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

AP ! TEVA PHARMS USA **500MG/10ML (50MG/ML)** **A040333 001** Jan 27, 2000

AP ! **2.5GM/50ML (50MG/ML)** **A040334 001** Feb 25, 2000

AP ! **5GM/100ML (50MG/ML)** **A040334 002** Feb 25, 2000

SOLUTION; TOPICAL

EFUDEX

AT +! BAUSCH **2%** **N016831 001**

AT +! **5%** **N016831 002**

FLUOROURACIL

AT TARO **2%** **A076526 001** Nov 05, 2003

AT **5%** **A076526 002** Nov 05, 2003

PRESCRIPTION DRUG PRODUCT LIST

FLUOXETINE HYDROCHLORIDE

CAPSULE;ORAL

FLUOXETINE HYDROCHLORIDE

<u>AB</u>	ALEMBIC PHARMS LTD	<u>EQ 40MG BASE</u>	<u>A090223 003</u>	Mar 19, 2009
<u>AB</u>	APNAR PHARMA LP	<u>EQ 40MG BASE</u>	<u>A075049 003</u>	Jan 29, 2002
<u>AB</u>	AUROBINDO PHARMA	<u>EQ 40MG BASE</u>	<u>A078619 003</u>	Jan 31, 2008
<u>AB</u>	CADILA PHARMS LTD	<u>EQ 40MG BASE</u>	<u>A206993 003</u>	May 23, 2019
<u>AB</u>	IVAX SUB TEVA PHARMS	<u>EQ 40MG BASE</u>	<u>A075245 003</u>	Sep 28, 2004
<u>AB</u>	MARKSANS PHARMA	<u>EQ 40MG BASE</u>	<u>A075465 003</u>	Aug 02, 2001
<u>AB</u>	SCIEGEN PHARMS INC	<u>EQ 40MG BASE</u>	<u>A204597 003</u>	Mar 16, 2015
<u>AB</u>	SUN PHARM INDS LTD	<u>EQ 40MG BASE</u>	<u>A076990 001</u>	Dec 13, 2004
<u>AB</u>	TEVA	<u>EQ 40MG BASE</u>	<u>A075452 003</u>	Jan 29, 2002

PROZAC

<u>AB</u>	<u>+</u> ELI LILLY AND CO	<u>EQ 40MG BASE</u>	<u>N018936 003</u>	Jun 15, 1999
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FLUOXETINE HYDROCHLORIDE

<u>AB1</u>	ALEMBIC PHARMS LTD	<u>EQ 10MG BASE</u>	<u>A090223 001</u>	Mar 19, 2009
<u>AB1</u>		<u>EQ 20MG BASE</u>	<u>A090223 002</u>	Mar 19, 2009
<u>AB1</u>	APNAR PHARMA LP	<u>EQ 10MG BASE</u>	<u>A075049 001</u>	Aug 02, 2001
<u>AB1</u>		<u>EQ 20MG BASE</u>	<u>A075049 002</u>	Jan 29, 2002
<u>AB1</u>	AUROBINDO PHARMA	<u>EQ 10MG BASE</u>	<u>A078619 001</u>	Jan 31, 2008
<u>AB1</u>		<u>EQ 20MG BASE</u>	<u>A078619 002</u>	Jan 31, 2008
<u>AB1</u>	BARR	<u>EQ 10MG BASE</u>	<u>A074803 002</u>	Jan 30, 2002
<u>AB1</u>		<u>EQ 20MG BASE</u>	<u>A074803 001</u>	Aug 02, 2001
<u>AB1</u>	CADILA PHARMS LTD	<u>EQ 10MG BASE</u>	<u>A206993 001</u>	May 23, 2019
<u>AB1</u>		<u>EQ 20MG BASE</u>	<u>A206993 002</u>	May 23, 2019
<u>AB1</u>	IVAX SUB TEVA PHARMS	<u>EQ 10MG BASE</u>	<u>A075245 002</u>	Jan 31, 2002
<u>AB1</u>		<u>EQ 20MG BASE</u>	<u>A075245 001</u>	Jan 31, 2002
<u>AB1</u>	LANDELA PHARM	<u>EQ 10MG BASE</u>	<u>A075464 001</u>	Jan 30, 2002
<u>AB1</u>		<u>EQ 20MG BASE</u>	<u>A075464 002</u>	Jan 30, 2002
<u>AB1</u>	MARKSANS PHARMA	<u>EQ 10MG BASE</u>	<u>A075465 001</u>	Jan 29, 2002
<u>AB1</u>		<u>EQ 20MG BASE</u>	<u>A075465 002</u>	Jan 29, 2002
<u>AB1</u>	SCIEGEN PHARMS INC	<u>EQ 10MG BASE</u>	<u>A204597 001</u>	Mar 16, 2015
<u>AB1</u>		<u>EQ 20MG BASE</u>	<u>A204597 002</u>	Mar 16, 2015
<u>AB1</u>	SPECGX LLC	<u>EQ 10MG BASE</u>	<u>A075658 001</u>	Jan 29, 2002
<u>AB1</u>		<u>EQ 20MG BASE</u>	<u>A075658 002</u>	Jan 29, 2002
<u>AB1</u>	TEVA	<u>EQ 10MG BASE</u>	<u>A075452 001</u>	Jan 29, 2002
<u>AB1</u>		<u>EQ 20MG BASE</u>	<u>A075452 002</u>	Jan 29, 2002
<u>AB1</u>	TEVA PHARMS USA	<u>EQ 10MG BASE</u>	<u>A076001 001</u>	Jan 29, 2002
<u>AB1</u>		<u>EQ 20MG BASE</u>	<u>A076001 002</u>	Jan 29, 2002

PROZAC

<u>AB1</u>	<u>+</u> ELI LILLY AND CO	<u>EQ 10MG BASE</u>	<u>N018936 006</u>	Dec 23, 1992
<u>AB1</u>	<u>+</u>	<u>EQ 20MG BASE</u>	<u>N018936 001</u>	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

<u>AA</u>	LANNETT CO INC	<u>EQ 20MG BASE/5ML</u>	<u>A077849 001</u>	Feb 09, 2007
<u>AA</u>	! PHARM ASSOC	<u>EQ 20MG BASE/5ML</u>	<u>A076015 001</u>	Jan 30, 2002
<u>AA</u>	SPECGX LLC	<u>EQ 20MG BASE/5ML</u>	<u>A075920 001</u>	Jan 29, 2002
<u>AA</u>	TEVA	<u>EQ 20MG BASE/5ML</u>	<u>A075506 001</u>	Aug 02, 2001
<u>AA</u>	WOCKHARDT BIO AG	<u>EQ 20MG BASE/5ML</u>	<u>A075514 001</u>	Aug 29, 2002

TABLET;ORAL

FLUOXETINE HYDROCHLORIDE

<u>AB</u>	ALEMBIC PHARMS LTD	<u>EQ 10MG BASE</u>	<u>A208698 001</u>	Apr 05, 2017
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A208698 002</u>	Apr 05, 2017
<u>AB</u>	<u>+</u> ! ALVOGEN	<u>EQ 60MG BASE</u>	<u>N202133 001</u>	Oct 06, 2011
<u>AB</u>	DR REDDYS LABS LTD	<u>EQ 10MG BASE</u>	<u>A076006 001</u>	Jan 30, 2002
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A076006 002</u>	Apr 23, 2018
<u>AB</u>		<u>EQ 60MG BASE</u>	<u>A211721 001</u>	Jan 25, 2019
<u>AB</u>	G AND W LABS INC	<u>EQ 60MG BASE</u>	<u>A212191 001</u>	Jul 05, 2019
<u>AB</u>	INVENTIA HLTHCARE	<u>EQ 60MG BASE</u>	<u>A209695 001</u>	Nov 20, 2017
<u>AB</u>	LUPIN LTD	<u>EQ 10MG BASE</u>	<u>A211653 001</u>	Apr 15, 2019
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A211653 002</u>	Apr 15, 2019
<u>AB</u>		<u>EQ 60MG BASE</u>	<u>A211632 001</u>	Feb 08, 2019
<u>AB</u>	MYLAN	<u>EQ 10MG BASE</u>	<u>A075755 001</u>	Aug 02, 2001
<u>AB</u>	!	<u>EQ 20MG BASE</u>	<u>A075755 002</u>	Aug 02, 2001
<u>AB</u>	PAR FORM	<u>EQ 10MG BASE</u>	<u>A203836 001</u>	Aug 19, 2016
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A203836 002</u>	Aug 19, 2016
<u>AB</u>	PAR PHARM INC	<u>EQ 60MG BASE</u>	<u>A209419 001</u>	Nov 16, 2017
<u>AB</u>	SCIEGEN PHARMS INC	<u>EQ 10MG BASE</u>	<u>A210935 001</u>	Mar 20, 2019

PRESCRIPTION DRUG PRODUCT LIST

FLUOXETINE HYDROCHLORIDE

TABLET; ORAL

FLUOXETINE HYDROCHLORIDE

<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A210935 002</u>	Mar 20, 2019
<u>AB</u>		<u>EQ 60MG BASE</u>	<u>A211282 001</u>	Jan 10, 2019
<u>AB</u>	TARO	<u>EQ 60MG BASE</u>	<u>A211477 001</u>	Nov 21, 2018
<u>AB</u>	TEVA	<u>EQ 10MG BASE</u>	<u>A075872 001</u>	Jan 29, 2002
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A075872 002</u>	Jan 04, 2019
<u>AB</u>	TEVA PHARMS USA	<u>EQ 60MG BASE</u>	<u>A211051 001</u>	Dec 03, 2018
<u>AB</u>	UPSHER SMITH LABS	<u>EQ 10MG BASE</u>	<u>A211696 001</u>	Jan 30, 2019
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A211696 002</u>	Jan 30, 2019
<u>AB1</u>	TORRENT	<u>EQ 10MG BASE</u>	<u>A206937 001</u>	Oct 21, 2016
<u>AB1</u>		<u>EQ 20MG BASE</u>	<u>A206937 002</u>	Oct 21, 2016
<u>SARAFEM</u>				
<u>AB1</u>	+ APIL	<u>EQ 10MG BASE</u>	<u>N021860 001</u>	May 19, 2006
<u>AB1</u>	+	<u>EQ 15MG BASE</u>	<u>N021860 002</u>	May 19, 2006
<u>AB1</u>	+	<u>EQ 20MG BASE</u>	<u>N021860 003</u>	May 19, 2006
<u>SELFEMRA</u>				
<u>AB1</u>	TEVA PHARMS USA	<u>EQ 10MG BASE</u>	<u>A200151 001</u>	Feb 03, 2014
<u>AB1</u>		<u>EQ 15MG BASE</u>	<u>A200151 002</u>	Feb 03, 2014
<u>AB1</u>		<u>EQ 20MG BASE</u>	<u>A200151 003</u>	Feb 03, 2014

FLUOXETINE HYDROCHLORIDE; OLANZAPINE

CAPSULE; ORAL

OLANZAPINE AND FLUOXETINE HYDROCHLORIDE

<u>AB</u>	PAR PHARM	<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>	SANDOZ	<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

FLUOXYMESTERONE

TABLET; ORAL

FLUOXYMESTERONE

! UPSHER SMITH LABS

10MG

A088342 001 Oct 21, 1983

FLUPHENAZINE DECANOATE

INJECTABLE; INJECTION

FLUPHENAZINE DECANOATE

<u>AO</u>	AUROBINDO PHARMA LTD	<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>	MYLAN LABS LTD	<u>25MG/ML</u>	<u>A075918 001</u>	Aug 17, 2001
<u>AO</u>	PAR STERILE PRODUCTS	<u>25MG/ML</u>	<u>A203732 001</u>	Jul 03, 2014
<u>AO</u>	WEST-WARD PHARMS INT	<u>25MG/ML</u>	<u>A074531 001</u>	Aug 30, 1996

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

PRESCRIPTION DRUG PRODUCT LIST

FLUPHENAZINE HYDROCHLORIDE

TABLET; ORAL

FLUPHENAZINE HYDROCHLORIDE

<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>	MYLAN	<u>1MG</u>	<u>A089804 002</u>	Aug 12, 1988
<u>AB</u>		<u>2.5MG</u>	<u>A089804 003</u>	Aug 12, 1988
<u>AB</u>		<u>5MG</u>	<u>A089804 004</u>	Aug 12, 1988
<u>AB</u>		<u>10MG</u>	<u>A089804 001</u>	Aug 12, 1988
<u>AB</u>	SANDOZ	<u>1MG</u>	<u>A089586 002</u>	Oct 16, 1987
<u>AB</u>		<u>2.5MG</u>	<u>A089586 003</u>	Oct 16, 1987
<u>AB</u>		<u>5MG</u>	<u>A089586 004</u>	Oct 16, 1987
<u>AB</u>		<u>10MG</u>	<u>A089586 001</u>	Oct 16, 1987

FLURANDRENOLIDE

CREAM; TOPICAL

CORDRAN SP

<u>AT</u>	+! ALMIRALL	<u>0.05%</u>	<u>N012806 002</u>	
<u>AT</u>	CINTEX SVCS	<u>0.05%</u>	<u>A205342 001</u>	Apr 13, 2016
	CORDRAN SP			
	+! ALMIRALL	0.025%	N012806 003	

LOTION; TOPICAL

CORDRAN

<u>AT</u>	+! ALMIRALL	<u>0.05%</u>	<u>N013790 001</u>	
<u>AT</u>	CINTEX SVCS	<u>0.05%</u>	<u>A205343 001</u>	Dec 22, 2016
<u>AT</u>	PERRIGO UK FINCO	<u>0.05%</u>	<u>A207133 001</u>	Aug 30, 2016

OINTMENT; TOPICAL

CORDRAN

<u>AT</u>	+! ALMIRALL	<u>0.05%</u>	<u>N012806 001</u>	
<u>AT</u>	TELLIGENT PHARMA INC	<u>0.05%</u>	<u>A207851 001</u>	Dec 30, 2016

TAPE; TOPICAL

CORDRAN

	+! ALMIRALL	0.004MG/SQ CM	N016455 001	
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FLURAZEPAM HYDROCHLORIDE

CAPSULE; ORAL

FLURAZEPAM HYDROCHLORIDE

	MYLAN PHARMS INC	15MG	A070345 002	Nov 27, 1985
	!	30MG	A070345 001	Nov 27, 1985

FLURBIPROFEN

TABLET; ORAL

FLURBIPROFEN

<u>AB</u>	! MYLAN	<u>100MG</u>	<u>A074358 002</u>	Jun 20, 1994
<u>AB</u>	TEVA	<u>100MG</u>	<u>A074431 001</u>	May 31, 1995
	MYLAN	50MG	A074358 001	Jun 20, 1994

FLURBIPROFEN SODIUM

SOLUTION/DROPS; OPHTHALMIC

FLURBIPROFEN SODIUM

<u>AT</u>	BAUSCH AND LOMB	<u>0.03%</u>	<u>A074447 001</u>	Jan 04, 1995
<u>AT</u>	+! ALLERGAN	<u>0.03%</u>	<u>N019404 001</u>	Dec 31, 1986

OCUFENFLUTAMIDE

CAPSULE; ORAL

FLUTAMIDE

<u>AB</u>	! CIPLA	<u>125MG</u>	<u>A075780 001</u>	Sep 19, 2001
<u>AB</u>	PAR PHARM	<u>125MG</u>	<u>A075298 001</u>	Sep 18, 2001

FLUTEMETAMOL F-18

INJECTABLE; INTRAVENOUS

VIZAMYL

	+! GE HEALTHCARE	121.5mCi/30ML (4.05mCi/ML)	N203137 002	Oct 25, 2013
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PRESCRIPTION DRUG PRODUCT LIST

FLUTICASONE FUROATE

POWDER; INHALATION

ARNUTTY 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

FLUTICASONE 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
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FLUTICASONE FUROATE; VILANTEROL TRIFENATATE

POWDER; INHALATION

BEO ELLIPTA

+	!	GLAXO GRP LTD	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

FLUTICASONE 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

FLUTICASONE PROPIONATE

AB		ACP NIMBLE	0.05%	A077055	001	Jun 30, 2006
AB		ANDA REPOSITORY	0.05%	A076633	001	May 14, 2004
AB		FOUGERA PHARMS	0.05%	A076451	001	May 14, 2004
AB	!	PERRIGO ISRAEL	0.05%	A076793	001	May 14, 2004

LOTION; TOPICAL

CUTIVATE

AB	+	!	FOUGERA PHARMS	0.05%	N021152	001	Mar 31, 2005
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FLUTICASONE PROPIONATE

AB		GLENMARK GENERICS	0.05%	A090759	001	May 02, 2011
AB		PERRIGO ISRAEL	0.05%	A091553	001	Jul 30, 2013

OINTMENT; TOPICAL

FLUTICASONE PROPIONATE

AB		ACP NIMBLE	0.005%	A077168	001	Mar 03, 2006
AB	!	PERRIGO NEW YORK	0.005%	A076668	001	May 14, 2004

POWDER; INHALATION

FLOVENT DISKUS 100

+	!	GLAXO GRP LTD	0.1MG/INH	N020833	002	Sep 29, 2000
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FLOVENT DISKUS 250

+	!	GLAXO GRP LTD	0.25MG/INH	N020833	003	Sep 29, 2000
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FLOVENT DISKUS 50

+	!	GLAXO GRP LTD	0.05MG/INH	N020833	001	Sep 29, 2000
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SPRAY, METERED; NASAL

FLUTICASONE PROPIONATE

AB		APOTEX INC	0.05MG/SPRAY	A077538	001	Sep 12, 2007
AB		HI TECH PHARMA	0.05MG/SPRAY	A077570	001	Jan 16, 2008
AB	!	HIKMA	0.05MG/SPRAY	A076504	001	Feb 22, 2006
AB		WOCKHARDT BIO AG	0.05MG/SPRAY	A078492	001	Jan 09, 2012

XHANCE

+	!	OPTINOSE US INC	0.093MG	N209022	001	Sep 18, 2017
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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	0.1MG/INH;EQ 0.05MG BASE/INH	N021077	001	Aug 24, 2000
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ADVAIR DISKUS 250/50

AB	+	!	GLAXO GRP LTD	0.25MG/INH;EQ 0.05MG BASE/INH	N021077	002	Aug 24, 2000
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ADVAIR DISKUS 500/50

AB	+	!	GLAXO GRP LTD	0.5MG/INH;EQ 0.05MG BASE/INH	N021077	003	Aug 24, 2000
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WIXELA INHUB

AB		MYLAN	0.1MG/INH;EQ 0.05MG BASE/INH	A208891	001	Jan 30, 2019
AB			0.25MG/INH;EQ 0.05MG BASE/INH	A208891	002	Jan 30, 2019
AB			0.5MG/INH;EQ 0.05MG BASE/INH	A208891	003	Jan 30, 2019

PRESCRIPTION DRUG PRODUCT LISTFLUTICASONE 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
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

<u>AB</u>	MYLAN PHARMS INC	<u>EQ 20MG BASE</u>	<u>A090595 001</u>	Apr 11, 2012
<u>AB</u>	!	<u>EQ 40MG BASE</u>	<u>A090595 002</u>	Apr 11, 2012
<u>AB</u>	TEVA PHARMS	<u>EQ 20MG BASE</u>	<u>A078407 001</u>	Jun 12, 2012
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A078407 002</u>	Jun 12, 2012

TABLET, EXTENDED RELEASE; ORAL

FLUVASTATIN SODIUM

<u>AB</u>	TEVA PHARMS USA	<u>EQ 80MG BASE</u>	<u>A079011 001</u>	Jan 27, 2016
<u>LESCOL XL</u>				
<u>AB</u>	+!	NOVARTIS	<u>EQ 80MG BASE</u>	<u>N021192 001</u>
				Oct 06, 2000

FLUVOXAMINE MALEATE

CAPSULE, EXTENDED RELEASE; ORAL

FLUVOXAMINE MALEATE

<u>AB</u>	ACTAVIS ELIZABETH	<u>100MG</u>	<u>A091482 001</u>	Apr 23, 2013
<u>AB</u>	!	<u>150MG</u>	<u>A091482 002</u>	Nov 18, 2013
<u>AB</u>	ANCHEN PHARMS	<u>100MG</u>	<u>A091476 001</u>	Mar 13, 2013
<u>AB</u>		<u>150MG</u>	<u>A091476 002</u>	Mar 13, 2013
<u>AB</u>	TORRENT	<u>100MG</u>	<u>A203240 001</u>	Oct 31, 2014
<u>AB</u>		<u>150MG</u>	<u>A203240 002</u>	Oct 31, 2014

TABLET; ORAL

FLUVOXAMINE MALEATE

<u>AB</u>	APOTEX	<u>25MG</u>	<u>A075902 001</u>	May 07, 2001
<u>AB</u>		<u>50MG</u>	<u>A075902 002</u>	May 07, 2001
<u>AB</u>		<u>100MG</u>	<u>A075902 003</u>	May 07, 2001
<u>AB</u>	TEVA	<u>25MG</u>	<u>A075893 001</u>	Sep 10, 2002
<u>AB</u>		<u>50MG</u>	<u>A075893 002</u>	Sep 10, 2002
<u>AB</u>		<u>100MG</u>	<u>A075893 003</u>	Sep 10, 2002
<u>AB</u>	UPSHER SMITH LABS	<u>25MG</u>	<u>A075888 001</u>	Nov 29, 2000
<u>AB</u>		<u>50MG</u>	<u>A075888 002</u>	Nov 29, 2000
<u>AB</u>	!	<u>100MG</u>	<u>A075888 003</u>	Nov 29, 2000
<u>LUVOX</u>				
<u>AB</u>	ANI PHARMS	<u>25MG</u>	<u>N021519 001</u>	Dec 20, 2007
<u>AB</u>		<u>50MG</u>	<u>N021519 002</u>	Dec 20, 2007
<u>AB</u>		<u>100MG</u>	<u>N021519 003</u>	Dec 20, 2007

FOLIC ACID

INJECTABLE; INJECTION

FOLIC ACID

<u>AP</u>	!	FRESENIUS KABI USA	<u>5MG/ML</u>	<u>A089202 001</u>	Feb 18, 1986
<u>AP</u>		XGEN PHARMS	<u>5MG/ML</u>	<u>A202522 001</u>	Nov 06, 2019

TABLET; ORAL

FOLIC ACID

<u>AA</u>	!	AMNEAL PHARM	<u>1MG</u>	<u>A040625 001</u>	Jul 21, 2005
<u>AA</u>		ATHEM	<u>1MG</u>	<u>A211064 001</u>	Mar 08, 2019
<u>AA</u>		CADILA PHARMS LTD	<u>1MG</u>	<u>A202437 001</u>	Jan 27, 2014
<u>AA</u>		CHARTWELL MOLECULAR	<u>1MG</u>	<u>A090035 001</u>	Jun 09, 2009
<u>AA</u>		HIKMA PHARMS	<u>1MG</u>	<u>A080600 001</u>	
<u>AA</u>		LEADING PHARMA LLC	<u>1MG</u>	<u>A040796 001</u>	Jan 12, 2009
<u>AA</u>		NUVO PHARMS INC	<u>1MG</u>	<u>A204418 001</u>	Jul 28, 2015
<u>AA</u>		QINGDAO BAHEAL PHARM	<u>1MG</u>	<u>A091145 001</u>	Jul 12, 2013
<u>AA</u>		VINTAGE	<u>1MG</u>	<u>A040756 001</u>	Jun 04, 2010
<u>AA</u>	!	WATSON LABS	<u>1MG</u>	<u>A080680 001</u>	

FOLLITROPIN ALFA/BETA

INJECTABLE; SUBCUTANEOUS

FOLLISTIM AQ

+	!	ORGANON USA INC	300 IU/0.36ML	N021211 001	Mar 23, 2004
+	!		600 IU/0.72ML	N021211 002	Mar 23, 2004
+	!		900 IU/1.08ML	N021211 004	Feb 11, 2005

PRESCRIPTION DRUG PRODUCT LIST

3-201 (of 453)

FOLLITROPIN ALFA/BETA

INJECTABLE; SUBCUTANEOUS

GONAL-F

+	!	EMD SERONO	450 IU/VIAL	N020378	005	Mar 26, 2004
+			1,050 IU/VIAL	N020378	004	Feb 28, 2001

GONAL-F RFF

+	!	EMD SERONO	75 IU/VIAL	N021765	002	Mar 25, 2004
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GONAL-F RFF RED1-JECT

+	!	EMD SERONO	150 IU/0.25ML	N021684	004	Nov 25, 2019
+			300 IU/0.5ML	N021684	001	May 25, 2004
+			450 IU/0.75ML	N021684	002	May 25, 2004
+			900 IU/1.5ML	N021684	003	May 25, 2004

FOMEPIZOLE

INJECTABLE; INJECTION

FOMEPIZOLE

AP		AM REGENT	<u>1.5GM/1.5ML (1GM/ML)</u>	<u>A078368</u>	<u>001</u>	Dec 14, 2007
AP	!	MYLAN INSTITUTIONAL	<u>1.5GM/1.5ML (1GM/ML)</u>	<u>A078639</u>	<u>001</u>	Mar 03, 2008
AP		NAVINTA LLC	<u>1.5GM/1.5ML (1GM/ML)</u>	<u>A078537</u>	<u>001</u>	Mar 06, 2008

FONDAPARINUX SODIUM

INJECTABLE; SUBCUTANEOUS

ARIXTRA

AP	+	!	MYLAN IRELAND LTD	<u>2.5MG/0.5ML</u>	<u>N021345</u>	<u>001</u>	Dec 07, 2001
AP	+			<u>5MG/0.4ML</u>	<u>N021345</u>	<u>002</u>	May 28, 2004
AP	+			<u>7.5MG/0.6ML</u>	<u>N021345</u>	<u>003</u>	May 28, 2004
AP	+			<u>10MG/0.8ML</u>	<u>N021345</u>	<u>004</u>	May 28, 2004

FONDAPARINUX SODIUM

AP		AUROBINDO PHARMA LTD	<u>2.5MG/0.5ML</u>	<u>A206918</u>	<u>001</u>	Dec 26, 2017
AP			<u>5MG/0.4ML</u>	<u>A206918</u>	<u>002</u>	Dec 26, 2017
AP			<u>7.5MG/0.6ML</u>	<u>A206918</u>	<u>003</u>	Dec 26, 2017
AP			<u>10MG/0.8ML</u>	<u>A206918</u>	<u>004</u>	Dec 26, 2017
AP		DR REDDYS LABS LTD	<u>2.5MG/0.5ML</u>	<u>A091316</u>	<u>001</u>	Jul 11, 2011
AP			<u>5MG/0.4ML</u>	<u>A091316</u>	<u>002</u>	Jul 11, 2011
AP			<u>7.5MG/0.6ML</u>	<u>A091316</u>	<u>003</u>	Jul 11, 2011
AP			<u>10MG/0.8ML</u>	<u>A091316</u>	<u>004</u>	Jul 11, 2011
AP		JIANGSU HENGRUI MED	<u>2.5MG/0.5ML</u>	<u>A206812</u>	<u>001</u>	May 15, 2018
AP			<u>5MG/0.4ML</u>	<u>A206812</u>	<u>002</u>	May 15, 2018
AP			<u>7.5MG/0.6ML</u>	<u>A206812</u>	<u>003</u>	May 15, 2018
AP			<u>10MG/0.8ML</u>	<u>A206812</u>	<u>004</u>	May 15, 2018
AP		SCINOPHARM TAIWAN	<u>2.5MG/0.5ML</u>	<u>A208615</u>	<u>001</u>	Nov 14, 2018
AP			<u>5MG/0.4ML</u>	<u>A208615</u>	<u>002</u>	Nov 14, 2018
AP			<u>7.5MG/0.6ML</u>	<u>A208615</u>	<u>003</u>	Nov 14, 2018
AP			<u>10MG/0.8ML</u>	<u>A208615</u>	<u>004</u>	Nov 14, 2018

FORMOTEROL FUMARATE

SOLUTION; INHALATION

PERFORMIST

+	!	MYLAN SPECTL	0.02MG/2ML	N022007	001	May 11, 2007
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FORMOTEROL FUMARATE; GLYCOPYRROLATE

AEROSOL, METERED; INHALATION

BEVESPI AEROSPHERE

+	!	ASTRAZENECA PHARMS	0.0048MG/INH;0.0090MG/INH	N208294	001	Apr 25, 2016
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FORMOTEROL FUMARATE; MOMETASONE FUROATE

AEROSOL, METERED; INHALATION

DULERA

+		MERCK SHARP DOHME	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

SUSPENSION; ORAL

LEXIVA

+	!	VIIV HLTHCARE	EQ 50MG BASE/ML	N022116	001	Jun 14, 2007
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TABLET; ORAL

FOSAMPRENAVIR CALCIUM

AB		MYLAN	<u>EQ 700MG BASE</u>	<u>A204060</u>	<u>001</u>	Apr 15, 2016	
AB		SUN PHARM	<u>EQ 700MG BASE</u>	<u>A204024</u>	<u>001</u>	Nov 20, 2019	
AB	+	!	VIIV HLTHCARE	<u>EQ 700MG BASE</u>	<u>N021548</u>	<u>001</u>	Oct 20, 2003

PRESCRIPTION DRUG PRODUCT LIST

FOSAPREPITANT DIMEGLUMINE

POWDER; INTRAVENOUS

EMEND

AP	+ !	MERCK AND CO INC	EQ 150MG BASE/VIAL	N022023 002	Nov 12, 2010
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FOSAPREPITANT DIMEGLUMINE

AP		APOTEX	EQ 150MG BASE/VIAL	A205020 001	Sep 05, 2019
AP		BAXTER HLTHCARE CORP	EQ 150MG BASE/VIAL	A211860 001	Sep 05, 2019
AP		BE PHARMS	EQ 150MG BASE/VIAL	A212309 001	Sep 05, 2019
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		SUNGEN PHARMA	EQ 150MG BASE/VIAL	A211624 001	Sep 05, 2019
		MYLAN LABS LTD	EQ 115MG BASE/VIAL	A204015 001	Sep 05, 2019
		TEVA PHARMS USA	EQ 150MG BASE/VIAL	N210064 001	Sep 05, 2019

FOSCARNET SODIUM

INJECTABLE; INJECTION

FOSCAVIR

+ !	CLINIGEN HLTHCARE	2.4GM/100ML	N020068 001	Sep 27, 1991
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FOSFOMYCIN TROMETHAMINE

FOR SOLUTION; ORAL

MONUROL

+ !	ZAMBON SPA	EQ 3GM BASE/PACKET	N050717 001	Dec 19, 1996
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FOSINOPRIL SODIUM

TABLET; ORAL

FOSINOPRIL SODIUM

AB		APOTEX INC	10MG	A076906 001	May 17, 2005
AB			20MG	A076906 002	May 17, 2005
AB			40MG	A076906 003	May 17, 2005
AB		AUROBINDO PHARMA LTD	10MG	A091163 001	Mar 30, 2011
AB			20MG	A091163 002	Mar 30, 2011
AB			40MG	A091163 003	Mar 30, 2011
AB		INVAGEN PHARMS	10MG	A077222 001	Apr 20, 2005
AB			20MG	A077222 002	Apr 20, 2005
AB			40MG	A077222 003	Apr 20, 2005
AB		PRINSTON INC	10MG	A205670 001	Aug 29, 2016
AB			20MG	A205670 002	Aug 29, 2016
AB			40MG	A205670 003	Aug 29, 2016
AB		TEVA	10MG	A076139 001	Nov 25, 2003
AB			20MG	A076139 002	Nov 25, 2003
AB	!		40MG	A076139 003	Nov 25, 2003
AB		UPSHER SMITH LABS	10MG	A076483 001	Apr 23, 2004
AB			20MG	A076483 002	Apr 23, 2004
AB			40MG	A076483 003	Apr 23, 2004

FOSINOPRIL SODIUM; HYDROCHLOROTHIAZIDE

TABLET; ORAL

FOSINOPRIL SODIUM AND HYDROCHLOROTHIAZIDE

AB		AUROBINDO PHARMA	10MG;12.5MG	A079245 001	Jul 09, 2009
AB			20MG;12.5MG	A079245 002	Jul 09, 2009
AB		EMCURE PHARMS LTD	10MG;12.5MG	A079025 001	Sep 17, 2010
AB	!		20MG;12.5MG	A079025 002	Sep 17, 2010
AB		INVAGEN PHARMS	10MG;12.5MG	A090228 001	Jul 09, 2009
AB			20MG;12.5MG	A090228 002	Jul 09, 2009
AB		SANDOZ	10MG;12.5MG	A076961 001	Sep 28, 2005
AB			20MG;12.5MG	A076961 002	Sep 28, 2005

FOSNETUPITANT CHLORIDE HYDROCHLORIDE; PALONOSETRON HYDROCHLORIDE

POWDER; INTRAVENOUS

AKYNZEO

+ !	HELSINN HLTHCARE	EQ 235MG BASE;EQ 0.25MG BASE	N210493 001	Apr 19, 2018
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FOSPHENYTOIN SODIUM

INJECTABLE; INJECTION

CEREBYX

AP	+ !	PARKE DAVIS	EQ 50MG PHENYTOIN NA/ML	N020450 001	Aug 05, 1996
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FOSPHENYTOIN SODIUM

AP		AMNEAL	EQ 50MG PHENYTOIN NA/ML	A078476 001	Mar 18, 2008
AP		FRESENIUS KABI USA	EQ 50MG PHENYTOIN NA/ML	A078052 001	Aug 06, 2007
AP		HIKMA FARMACEUTICA	EQ 50MG PHENYTOIN NA/ML	A078765 001	Dec 02, 2009
AP		MYLAN LABS LTD	EQ 50MG PHENYTOIN NA/ML	A078736 001	Jun 08, 2010

PRESCRIPTION DRUG PRODUCT LIST

FOSPHENYTOIN SODIUM

INJECTABLE; INJECTION

FOSPHENYTOIN SODIUM

<u>AP</u>	SUN PHARM	<u>EQ 50MG PHENYTOIN NA/ML</u>	<u>A078417 001</u>	Mar 18, 2008
<u>AP</u>	WEST-WARD PHARMS INT	<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>	WOCKHARDT	<u>EQ 50MG PHENYTOIN NA/ML</u>	<u>A078137 001</u>	Aug 06, 2007

FOSTAMATINIB DISODIUM

TABLET; ORAL

TAVALISSE

+	RIGEL PHARMS INC	EQ 100MG BASE	N209299 001	Apr 17, 2018
+	!	EQ 150MG BASE	N209299 002	Apr 17, 2018

FROVATRIPTAN SUCCINATE

TABLET; ORAL

FROVA

<u>AB</u>	+	ENDO PHARMS	<u>EQ 2.5MG BASE</u>	<u>N021006 001</u>	Nov 08, 2001
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FROVATRIPTAN SUCCINATE

<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>		MYLAN	<u>EQ 2.5MG BASE</u>	<u>A202931 001</u>	Aug 28, 2014

FULVESTRANT

INJECTABLE; INTRAMUSCULAR

FASLODEX

<u>AO</u>	+	ASTRAZENECA	<u>50MG/ML</u>	<u>N021344 001</u>	Apr 25, 2002
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FULVESTRANT

<u>AO</u>		AMNEAL	<u>50MG/ML</u>	<u>A210044 001</u>	Mar 04, 2019
<u>AO</u>		CHIA TAI TIANQING	<u>50MG/ML</u>	<u>A211422 001</u>	Feb 07, 2020
<u>AO</u>		GLENMARK PHARMS INC	<u>50MG/ML</u>	<u>A207754 001</u>	Aug 22, 2019
<u>AO</u>		HBT LABS INC	<u>50MG/ML</u>	<u>A209714 001</u>	Nov 21, 2019
<u>AO</u>		MYLAN INSTITUTIONAL	<u>50MG/ML</u>	<u>A208811 001</u>	Jul 23, 2019
<u>AO</u>		SAGENT PHARMS INC	<u>50MG/ML</u>	<u>A205871 001</u>	Aug 22, 2019
<u>AO</u>		SANDOZ INC	<u>50MG/ML</u>	<u>A205935 001</u>	May 14, 2019

SOLUTION; INTRAMUSCULAR

FULVESTRANT

	FRESENIUS KABI USA	250MG/5ML (50MG/ML)	N210326 001	May 20, 2019
	TEVA PHARMS USA INC	250MG/5ML (50MG/ML)	N210063 001	Aug 19, 2019

FUROSEMIDE

INJECTABLE; INJECTION

FUROSEMIDE

<u>AP</u>		AMNEAL PHARMS CO	<u>10MG/ML</u>	<u>A207552 001</u>	Jul 20, 2016
<u>AP</u>		AUROBINDO PHARMA LTD	<u>10MG/ML</u>	<u>A212174 001</u>	May 03, 2019
<u>AP</u>	!	BAXTER HLTHCARE CORP	<u>10MG/ML</u>	<u>A202747 001</u>	Jan 27, 2014
<u>AP</u>		EMCURE PHARMS LTD	<u>10MG/ML</u>	<u>A203428 001</u>	Aug 26, 2014
<u>AP</u>		FRESENIUS KABI USA	<u>10MG/ML</u>	<u>N018902 001</u>	May 22, 1984
<u>AP</u>		HOSPIRA	<u>10MG/ML</u>	<u>A075241 001</u>	May 28, 1999
<u>AP</u>			<u>10MG/ML</u>	<u>N018667 001</u>	May 28, 1982
<u>AP</u>		WOCKHARDT	<u>10MG/ML</u>	<u>A077941 001</u>	Mar 22, 2007

SOLUTION; ORAL

FUROSEMIDE

<u>AA</u>	!	HIKMA	<u>10MG/ML</u>	<u>A070434 001</u>	Apr 22, 1987
<u>AA</u>		WOCKHARDT BIO AG	<u>10MG/ML</u>	<u>A070655 001</u>	Oct 02, 1987
		HIKMA	40MG/5ML	A070433 001	Apr 22, 1987

TABLET; ORAL

FUROSEMIDE

<u>AB</u>		HIKMA	<u>20MG</u>	<u>N018823 001</u>	Nov 10, 1983
<u>AB</u>			<u>40MG</u>	<u>N018823 002</u>	Nov 10, 1983
<u>AB</u>			<u>80MG</u>	<u>A070086 001</u>	Jan 24, 1986
<u>AB</u>		IPCA LABS LTD	<u>20MG</u>	<u>A078010 001</u>	Sep 18, 2006
<u>AB</u>			<u>40MG</u>	<u>A078010 002</u>	Sep 18, 2006
<u>AB</u>			<u>80MG</u>	<u>A078010 003</u>	Sep 18, 2006
<u>AB</u>		LEADING PHARMA LLC	<u>20MG</u>	<u>A077293 001</u>	Nov 09, 2005
<u>AB</u>			<u>40MG</u>	<u>A077293 002</u>	Nov 09, 2005
<u>AB</u>			<u>80MG</u>	<u>A077293 003</u>	Nov 09, 2005
<u>AB</u>		MYLAN	<u>20MG</u>	<u>N018487 001</u>	
<u>AB</u>			<u>40MG</u>	<u>N018487 002</u>	
<u>AB</u>			<u>80MG</u>	<u>A070082 001</u>	Oct 29, 1986
<u>AB</u>		PRINSTON INC	<u>20MG</u>	<u>A076796 001</u>	Mar 26, 2004
<u>AB</u>			<u>40MG</u>	<u>A076796 002</u>	Mar 26, 2004

PRESCRIPTION DRUG PRODUCT LISTFUROSEMIDE

TABLET; ORAL

FUROSEMIDE

<u>AB</u>		<u>80MG</u>	<u>A076796 003</u>	Mar 26, 2004
<u>AB</u>	SANDOZ	<u>20MG</u>	<u>N018569 002</u>	
<u>AB</u>		<u>40MG</u>	<u>N018569 001</u>	
<u>AB</u>		<u>80MG</u>	<u>N018569 005</u>	Aug 14, 1984
<u>LASIX</u>				
<u>AB</u>	+	US PHARM HOLDINGS	<u>20MG</u>	<u>N016273 002</u>
<u>AB</u>	+		<u>40MG</u>	<u>N016273 001</u>
<u>AB</u>	+		<u>80MG</u>	<u>N016273 003</u>

GABAPENTIN

CAPSULE; ORAL

GABAPENTIN

<u>AB</u>	ACI HEALTHCARE LTD	<u>100MG</u>	<u>A206943 001</u>	May 14, 2018
<u>AB</u>		<u>300MG</u>	<u>A206943 002</u>	May 14, 2018
<u>AB</u>		<u>400MG</u>	<u>A206943 003</u>	May 14, 2018
<u>AB</u>	ACTAVIS ELIZABETH	<u>100MG</u>	<u>A075350 001</u>	Sep 12, 2003
<u>AB</u>		<u>300MG</u>	<u>A075350 002</u>	Sep 12, 2003
<u>AB</u>		<u>400MG</u>	<u>A075350 003</u>	Sep 12, 2003
<u>AB</u>	ALKEM	<u>100MG</u>	<u>A090858 001</u>	Dec 17, 2010
<u>AB</u>		<u>300MG</u>	<u>A090858 002</u>	Dec 17, 2010
<u>AB</u>		<u>400MG</u>	<u>A090858 003</u>	Dec 17, 2010
<u>AB</u>	AMNEAL PHARMS NY	<u>100MG</u>	<u>A078428 001</u>	Jul 25, 2007
<u>AB</u>		<u>300MG</u>	<u>A078428 002</u>	Jul 25, 2007
<u>AB</u>		<u>400MG</u>	<u>A078428 003</u>	Jul 25, 2007
<u>AB</u>	AUROBINDO PHARMA LTD	<u>100MG</u>	<u>A078787 001</u>	Jan 31, 2008
<u>AB</u>		<u>300MG</u>	<u>A078787 002</u>	Jan 31, 2008
<u>AB</u>		<u>400MG</u>	<u>A078787 003</u>	Jan 31, 2008
<u>AB</u>	CSPC OUYI	<u>100MG</u>	<u>A075477 001</u>	Mar 23, 2005
<u>AB</u>		<u>300MG</u>	<u>A075477 002</u>	Mar 23, 2005
<u>AB</u>		<u>400MG</u>	<u>A075477 003</u>	Mar 23, 2005
<u>AB</u>	EPIC PHARMA LLC	<u>100MG</u>	<u>A207099 001</u>	Mar 24, 2017
<u>AB</u>		<u>300MG</u>	<u>A207099 002</u>	Mar 24, 2017
<u>AB</u>		<u>400MG</u>	<u>A207099 003</u>	Mar 24, 2017
<u>AB</u>	GRANULES INDIA LTD	<u>100MG</u>	<u>A075360 001</u>	Apr 06, 2005
<u>AB</u>		<u>300MG</u>	<u>A075360 002</u>	Apr 06, 2005
<u>AB</u>		<u>400MG</u>	<u>A075360 003</u>	Apr 06, 2005
<u>AB</u>	INVAGEN PHARMS	<u>100MG</u>	<u>A090705 001</u>	Dec 30, 2009
<u>AB</u>		<u>300MG</u>	<u>A090705 002</u>	Dec 30, 2009
<u>AB</u>		<u>400MG</u>	<u>A090705 003</u>	Dec 30, 2009
<u>AB</u>	JIANGSU HENGRUI MED	<u>100MG</u>	<u>A091008 001</u>	Oct 26, 2017
<u>AB</u>		<u>300MG</u>	<u>A091008 002</u>	Oct 26, 2017
<u>AB</u>		<u>400MG</u>	<u>A091008 003</u>	Oct 26, 2017
<u>AB</u>	MARKSANS PHARMA	<u>100MG</u>	<u>A090007 001</u>	Jul 21, 2011
<u>AB</u>		<u>300MG</u>	<u>A090007 002</u>	Jul 21, 2011
<u>AB</u>		<u>400MG</u>	<u>A090007 003</u>	Jul 21, 2011
<u>AB</u>	MYLAN	<u>100MG</u>	<u>A090158 001</u>	Feb 14, 2011
<u>AB</u>		<u>300MG</u>	<u>A090158 002</u>	Feb 14, 2011
<u>AB</u>		<u>400MG</u>	<u>A090158 003</u>	Feb 14, 2011
<u>AB</u>	SCIEGEN PHARMS INC	<u>100MG</u>	<u>A204989 001</u>	Feb 18, 2016
<u>AB</u>		<u>300MG</u>	<u>A204989 002</u>	Feb 18, 2016
<u>AB</u>		<u>400MG</u>	<u>A204989 003</u>	Feb 18, 2016
<u>AB</u>	STRIDES PHARMA	<u>100MG</u>	<u>A211314 001</u>	Oct 16, 2018
<u>AB</u>		<u>300MG</u>	<u>A211314 002</u>	Oct 16, 2018
<u>AB</u>		<u>400MG</u>	<u>A211314 003</u>	Oct 16, 2018
<u>AB</u>	SUN PHARM INDS LTD	<u>100MG</u>	<u>A077242 001</u>	Aug 24, 2006
<u>AB</u>		<u>300MG</u>	<u>A077242 002</u>	Aug 24, 2006
<u>AB</u>		<u>400MG</u>	<u>A077242 003</u>	Aug 24, 2006
<u>AB</u>	TARO	<u>100MG</u>	<u>A077261 001</u>	Aug 02, 2013
<u>AB</u>		<u>300MG</u>	<u>A077261 002</u>	Aug 02, 2013
<u>AB</u>		<u>400MG</u>	<u>A077261 003</u>	Aug 02, 2013
<u>AB</u>	TEVA PHARMS	<u>100MG</u>	<u>A075435 001</u>	Oct 08, 2004
<u>AB</u>		<u>300MG</u>	<u>A075435 002</u>	Oct 08, 2004
<u>AB</u>		<u>400MG</u>	<u>A075435 003</u>	Oct 08, 2004
<u>NEURONTIN</u>				
<u>AB</u>	+	PFIZER PHARMS	<u>100MG</u>	<u>N020235 001</u> Dec 30, 1993
<u>AB</u>	+		<u>300MG</u>	<u>N020235 002</u> Dec 30, 1993
<u>AB</u>	+		<u>400MG</u>	<u>N020235 003</u> Dec 30, 1993

PRESCRIPTION DRUG PRODUCT LIST

GABAPENTIN

SOLUTION;ORAL

GABAPENTIN

<u>AA</u>	ACELLA PHARMS LLC	<u>250MG/5ML</u>	<u>A076403</u>	<u>001</u>	May 01, 2012
<u>AA</u>	AMNEAL PHARMS	<u>250MG/5ML</u>	<u>A202024</u>	<u>001</u>	Mar 23, 2012
<u>AA</u>	HI TECH PHARMA	<u>250MG/5ML</u>	<u>A078974</u>	<u>001</u>	Feb 18, 2011
<u>AA</u>	TARO	<u>250MG/5ML</u>	<u>A076672</u>	<u>001</u>	Jul 03, 2013
<u>AA</u>	TRIS PHARMA INC	<u>250MG/5ML</u>	<u>A091286</u>	<u>001</u>	Mar 14, 2016
<u>AA</u>	VISTAPHARM	<u>250MG/5ML</u>	<u>A211330</u>	<u>001</u>	Dec 03, 2019

NEURONTIN

<u>AA</u>	<u>+</u> PARKE DAVIS	<u>250MG/5ML</u>	<u>N021129</u>	<u>001</u>	Mar 02, 2000
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TABLET;ORAL

GABAPENTIN

<u>AB</u>	ACI HEALTHCARE LTD	<u>600MG</u>	<u>A203244</u>	<u>002</u>	Jul 12, 2013
<u>AB</u>		<u>800MG</u>	<u>A203244</u>	<u>001</u>	Jul 12, 2013
<u>AB</u>	ACTAVIS ELIZABETH	<u>600MG</u>	<u>A075694</u>	<u>001</u>	Oct 21, 2004
<u>AB</u>		<u>800MG</u>	<u>A075694</u>	<u>002</u>	Oct 21, 2004
<u>AB</u>	ALKEM LABS LTD	<u>600MG</u>	<u>A206402</u>	<u>001</u>	Dec 23, 2015
<u>AB</u>		<u>800MG</u>	<u>A206402</u>	<u>002</u>	Dec 23, 2015
<u>AB</u>	APOTEX INC	<u>100MG</u>	<u>A077894</u>	<u>001</u>	Oct 10, 2006
<u>AB</u>		<u>300MG</u>	<u>A077894</u>	<u>002</u>	Oct 10, 2006
<u>AB</u>		<u>400MG</u>	<u>A077894</u>	<u>003</u>	Oct 10, 2006
<u>AB</u>	AUROBINDO PHARMA LTD	<u>600MG</u>	<u>A200651</u>	<u>001</u>	Oct 06, 2011
<u>AB</u>		<u>800MG</u>	<u>A200651</u>	<u>002</u>	Oct 06, 2011
<u>AB</u>	CSPC OUYI	<u>600MG</u>	<u>A207057</u>	<u>001</u>	Oct 26, 2017
<u>AB</u>		<u>800MG</u>	<u>A207057</u>	<u>002</u>	Oct 26, 2017
<u>AB</u>	GLENMARK PHARMS LTD	<u>600MG</u>	<u>A077662</u>	<u>001</u>	Aug 18, 2006
<u>AB</u>		<u>800MG</u>	<u>A077662</u>	<u>002</u>	Aug 18, 2006
<u>AB</u>	INVAGEN PHARMS	<u>600MG</u>	<u>A202764</u>	<u>001</u>	Oct 16, 2012
<u>AB</u>		<u>800MG</u>	<u>A202764</u>	<u>002</u>	Oct 16, 2012
<u>AB</u>	IVAX SUB TEVA PHARMS	<u>100MG</u>	<u>A076017</u>	<u>001</u>	Apr 28, 2004
<u>AB</u>		<u>300MG</u>	<u>A076017</u>	<u>002</u>	Apr 28, 2004
<u>AB</u>		<u>400MG</u>	<u>A076017</u>	<u>003</u>	Apr 28, 2004
<u>AB</u>	MYLAN PHARMS INC	<u>600MG</u>	<u>A090335</u>	<u>001</u>	Jun 01, 2010
<u>AB</u>		<u>800MG</u>	<u>A090335</u>	<u>002</u>	Jun 01, 2010
<u>AB</u>	RUBICON	<u>600MG</u>	<u>A077661</u>	<u>004</u>	Sep 13, 2006
<u>AB</u>		<u>800MG</u>	<u>A077661</u>	<u>005</u>	Sep 13, 2006
<u>AB</u>	SCIEGEN PHARMS INC	<u>600MG</u>	<u>A205101</u>	<u>001</u>	Feb 04, 2016
<u>AB</u>		<u>800MG</u>	<u>A205101</u>	<u>002</u>	Feb 04, 2016
<u>AB</u>	SUN PHARM INDS LTD	<u>600MG</u>	<u>A077525</u>	<u>001</u>	Aug 24, 2006
<u>AB</u>		<u>800MG</u>	<u>A077525</u>	<u>002</u>	Aug 24, 2006
<u>AB</u>	TEVA PHARMS USA	<u>600MG</u>	<u>A205807</u>	<u>001</u>	Mar 10, 2017
<u>AB</u>		<u>800MG</u>	<u>A205807</u>	<u>002</u>	Mar 10, 2017
<u>AB</u>	ZYDUS PHARMS USA INC	<u>600MG</u>	<u>A078926</u>	<u>001</u>	Feb 11, 2011
<u>AB</u>		<u>800MG</u>	<u>A078926</u>	<u>002</u>	Feb 11, 2011

NEURONTIN

<u>AB</u>	<u>+</u> PFIZER PHARMS	<u>600MG</u>	<u>N020882</u>	<u>001</u>	Oct 09, 1998
<u>AB</u>	<u>+</u>	<u>800MG</u>	<u>N020882</u>	<u>002</u>	Oct 09, 1998

GRALISE

BX	<u>+</u> ALMATICA	300MG	N022544	001	Jan 28, 2011
BX	<u>+</u>	600MG	N022544	002	Jan 28, 2011

GABAPENTIN ENACARBIL

TABLET, EXTENDED RELEASE;ORAL

HORIZANT

<u>+</u>	ARBOR PHARMS LLC	300MG	N022399	002	Dec 13, 2011
<u>+</u>		600MG	N022399	001	Apr 06, 2011

GADOBENATE DIMEGLUMINE

INJECTABLE; INTRAVENOUS

MULTIHANCE

<u>+</u>	BRACCO	2.645GM/5ML (529MG/ML)	N021357	001	Nov 23, 2004
<u>+</u>		5.29GM/10ML (529MG/ML)	N021357	002	Nov 23, 2004
<u>+</u>		7.935GM/15ML (529MG/ML)	N021357	003	Nov 23, 2004
<u>+</u>		10.58GM/20ML (529MG/ML)	N021357	004	Nov 23, 2004

MULTIHANCE MULTIPACK

<u>+</u>	BRACCO	26.45GM/50ML (529MG/ML)	N021358	001	Nov 23, 2004
<u>+</u>		52.9GM/100ML (529MG/ML)	N021358	002	Nov 23, 2004

PRESCRIPTION DRUG PRODUCT LIST

GADOBUTROL

SOLUTION; INTRAVENOUS

GADAVIST

+	!	BAYER HLTHCARE	1.20944GM/2ML (604.72MG/ML)	N201277	006	Dec 18, 2013
+	!		4.5354GM/7.5ML (604.72MG/ML)	N201277	001	Mar 14, 2011
+	!		6.0472GM/10ML (604.72MG/ML)	N201277	002	Mar 14, 2011
+	!		9.0708GM/15ML (604.72MG/ML)	N201277	003	Mar 14, 2011
+	!		18.1416GM/30ML (604.72MG/ML)	N201277	004	Mar 14, 2011
+	!		39.3068GM/65ML (604.72MG/ML)	N201277	005	Mar 14, 2011

GADODIAMIDE

INJECTABLE; INJECTION

OMNISCAN

+	!	GE HEALTHCARE	287MG/ML	N020123	001	Jan 08, 1993
+	!		28.7GM/100ML (287MG/ML)	N022066	002	Sep 05, 2007

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>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

DOTAREM

<u>AP</u>	+	!	GUERBET	<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
	+	!		37.69GM/100ML (376.9MG/ML)	N204781	001	Mar 20, 2013
	+	!		1.8845GM/5ML (376.9MG/ML)	N204781	005	Mar 31, 2017

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 LTD	<u>EQ 8MG BASE</u>	<u>A204895</u>	<u>001</u>	Aug 05, 2016
<u>AB</u>			<u>EQ 16MG BASE</u>	<u>A204895</u>	<u>002</u>	Aug 05, 2016
<u>AB</u>			<u>EQ 24MG BASE</u>	<u>A204895</u>	<u>003</u>	Aug 05, 2016
<u>AB</u>		BARR	<u>EQ 8MG BASE</u>	<u>A078189</u>	<u>001</u>	Sep 15, 2008
<u>AB</u>			<u>EQ 16MG BASE</u>	<u>A078189</u>	<u>002</u>	Sep 15, 2008
<u>AB</u>			<u>EQ 24MG BASE</u>	<u>A078189</u>	<u>003</u>	Sep 15, 2008
<u>AB</u>		SUN PHARM	<u>EQ 8MG BASE</u>	<u>A090178</u>	<u>001</u>	Feb 02, 2011
<u>AB</u>			<u>EQ 16MG BASE</u>	<u>A090178</u>	<u>002</u>	Feb 02, 2011
<u>AB</u>			<u>EQ 24MG BASE</u>	<u>A090178</u>	<u>003</u>	Feb 02, 2011
<u>AB</u>		WATSON LABS	<u>EQ 8MG BASE</u>	<u>A079028</u>	<u>001</u>	Dec 15, 2008
<u>AB</u>			<u>EQ 16MG BASE</u>	<u>A079028</u>	<u>002</u>	Dec 15, 2008
<u>AB</u>			<u>EQ 24MG BASE</u>	<u>A079028</u>	<u>003</u>	Dec 15, 2008

RAZADYNE ER

<u>AB</u>	+	!	JANSSEN PHARMS	<u>EQ 8MG BASE</u>	<u>N021615</u>	<u>001</u>	Apr 01, 2005
<u>AB</u>	+			<u>EQ 16MG BASE</u>	<u>N021615</u>	<u>002</u>	Apr 01, 2005
<u>AB</u>	+			<u>EQ 24MG BASE</u>	<u>N021615</u>	<u>003</u>	Apr 01, 2005

SOLUTION; ORAL

GALANTAMINE HYDROBROMIDE

!		HIKMA	4MG/ML	A078185	001	Jan 30, 2009
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TABLET; ORAL

GALANTAMINE HYDROBROMIDE

<u>AB</u>		APOTEX INC	<u>EQ 4MG BASE</u>	<u>A077781</u>	<u>001</u>	Sep 27, 2011
<u>AB</u>			<u>EQ 8MG BASE</u>	<u>A077781</u>	<u>002</u>	Sep 27, 2011
<u>AB</u>			<u>EQ 12MG BASE</u>	<u>A077781</u>	<u>003</u>	Sep 27, 2011
<u>AB</u>		AUROBINDO PHARMA LTD	<u>EQ 4MG BASE</u>	<u>A090957</u>	<u>001</u>	Mar 29, 2011
<u>AB</u>			<u>EQ 8MG BASE</u>	<u>A090957</u>	<u>002</u>	Mar 29, 2011
<u>AB</u>			<u>EQ 12MG BASE</u>	<u>A090957</u>	<u>003</u>	Mar 29, 2011
<u>AB</u>		BARR	<u>EQ 4MG BASE</u>	<u>A077605</u>	<u>001</u>	Aug 28, 2008

PRESCRIPTION DRUG PRODUCT LIST

3-207 (of 453)

GALANTAMINE HYDROBROMIDE

TABLET; ORAL

GALANTAMINE HYDROBROMIDE

<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
<u>RAZADYNE</u>				
<u>AB</u>	+! JANSSEN PHARMS	<u>EQ 4MG BASE</u>	<u>N021169 001</u>	Feb 28, 2001
<u>AB</u>	+	<u>EQ 8MG BASE</u>	<u>N021169 002</u>	Feb 28, 2001
<u>AB</u>	+	<u>EQ 12MG BASE</u>	<u>N021169 003</u>	Feb 28, 2001

GALLIUM CITRATE GA-67

INJECTABLE; INJECTION

GALLIUM CITRATE GA 67

BS	CURIUM	2mCi/ML	N018058 001	
BS	LANTHEUS MEDCL	2mCi/ML	N017478 001	

GALLIUM DOTATATE GA-68

POWDER; INTRAVENOUS

NETSPOT

	+! AAA USA INC	2.1-5.5mCi/ML	N208547 001	Jun 01, 2016
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GALLIUM DOTATOC GA-68

SOLUTION; INTRAVENOUS

GALLIUM DOTATOC GA 68

	+! UIHC PET IMAGING	0.5-4mCi/ML	N210828 001	Aug 21, 2019
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GANCICLOVIR

GEL; OPHTHALMIC

ZIRGAN

	+! BAUSCH AND LOMB	0.15%	N022211 001	Sep 15, 2009
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SOLUTION; INTRAVENOUS

GANZYK-RTU

	+! EXELA PHARMA SCS LLC	500MG/250ML (2MG/ML)	N209347 001	Feb 17, 2017
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GANCICLOVIR SODIUM

INJECTABLE; INJECTION

CYTOVENE

<u>AP</u>	+! CHEPLAPHARM	<u>EQ 500MG BASE/VIAL</u>	<u>N019661 001</u>	Jun 23, 1989
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GANCICLOVIR SODIUM

<u>AP</u>	FRESENIUS KABI USA	<u>EQ 500MG BASE/VIAL</u>	<u>A090658 001</u>	Jun 21, 2010
<u>AP</u>	HAINAN POLY PHARM	<u>EQ 500MG BASE/VIAL</u>	<u>A204204 001</u>	Nov 08, 2018
<u>AP</u>	MYLAN LABS LTD	<u>EQ 500MG BASE/VIAL</u>	<u>A204560 001</u>	Nov 17, 2017
<u>AP</u>	PAR STERILE PRODUCTS	<u>EQ 500MG BASE/VIAL</u>	<u>A204950 001</u>	Dec 06, 2016
<u>AP</u>	PHARMASCIENCE INC	<u>EQ 500MG BASE/VIAL</u>	<u>A207645 001</u>	Dec 08, 2017

GANIRELIX ACETATE

INJECTABLE; INJECTION

GANIRELIX ACETATE

<u>AP</u>	+! ORGANON USA INC	<u>250MCG/0.5ML</u>	<u>N021057 001</u>	Jul 29, 1999
<u>AP</u>	SUN PHARM	<u>250MCG/0.5ML</u>	<u>A204246 001</u>	Nov 30, 2018

GATIFLOXACIN

SOLUTION/DROPS; OPHTHALMIC

GATIFLOXACIN

<u>AT</u>	HI-TECH PHARMA CO	<u>0.5%</u>	<u>A203189 001</u>	Sep 03, 2014
<u>AT</u>	LUPIN LTD	<u>0.5%</u>	<u>A202653 001</u>	Aug 28, 2013
<u>AT</u>	MYLAN	<u>0.5%</u>	<u>A206446 001</u>	Jun 08, 2018
<u>AT</u>	SANDOZ INC	<u>0.5%</u>	<u>A204227 001</u>	Jul 11, 2016

ZYMAXID

<u>AT</u>	+! ALLERGAN	<u>0.5%</u>	<u>N022548 001</u>	May 18, 2010
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PRESCRIPTION DRUG PRODUCT LIST

3-208 (of 453)

GATIFLOXACIN

SOLUTION/DROPS;OPHTHALMIC

ZYMAR

+! ALLERGAN 0.3% N021493 001 Mar 28, 2003

GEFITINIB

TABLET;ORAL

IRESSA

+! ASTRAZENECA PHARMS 250MG N206995 001 Jul 13, 2015

GEMCITABINE HYDROCHLORIDE

INJECTABLE; INJECTION

GEMCITABINE HYDROCHLORIDE

<u>AP</u>	ACCORD HLTHCARE	<u>EQ 200MG BASE/VIAL</u>	<u>A091594 001</u>	Jul 25, 2011
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>A091594 002</u>	Jul 25, 2011
<u>AP</u>		<u>EQ 2GM BASE/VIAL</u>	<u>A091594 003</u>	Jul 25, 2011
<u>AP</u>	CIPLA	<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
<u>AP</u>	DR REDDYS LABS LTD	<u>EQ 200MG BASE/VIAL</u>	<u>A091365 001</u>	Jul 25, 2011
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>A091365 002</u>	Jul 25, 2011
<u>AP</u>		<u>EQ 2GM BASE/VIAL</u>	<u>A202997 001</u>	May 07, 2013
<u>AP</u>	EMCURE PHARMS LTD	<u>EQ 200MG BASE/VIAL</u>	<u>A202063 001</u>	Sep 11, 2012
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>A202063 002</u>	Sep 11, 2012
<u>AP</u>	FRESENIUS KABI USA	<u>EQ 200MG BASE/VIAL</u>	<u>A090799 001</u>	Jul 25, 2011
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>A090799 002</u>	Jul 25, 2011
<u>AP</u>		<u>EQ 2GM BASE/VIAL</u>	<u>A090242 003</u>	May 16, 2011
<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>	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>	JIANGSU HANSOH PHARM	<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>	MYLAN LABS LTD	<u>EQ 200MG BASE/VIAL</u>	<u>A200145 001</u>	Jul 25, 2011
<u>AP</u>		<u>200MG/5.26ML (38MG/ML)</u>	<u>A205242 001</u>	Dec 06, 2017
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>A200145 002</u>	Jul 25, 2011
<u>AP</u>		<u>1GM/26.3ML (38MG/ML)</u>	<u>A205242 002</u>	Dec 06, 2017
<u>AP</u>		<u>EQ 2GM BASE/VIAL</u>	<u>A200145 003</u>	Jul 25, 2011
<u>AP</u>		<u>2GM/52.6ML (38MG/ML)</u>	<u>A205242 003</u>	Dec 06, 2017
<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 MEDICARE LTD	<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>	TEVA PHARMS	<u>EQ 1GM BASE/VIAL</u>	<u>A077983 001</u>	Jan 25, 2011
<u>AP</u>		<u>EQ 200MG BASE/VIAL</u>	<u>A077983 002</u>	Jan 25, 2011

GEMZAR

<u>AP</u>	+! LILLY	<u>EQ 200MG BASE/VIAL</u>	<u>N020509 001</u>	May 15, 1996
<u>AP</u>	+!	<u>EQ 1GM BASE/VIAL</u>	<u>N020509 002</u>	May 15, 1996

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

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

PRESCRIPTION DRUG PRODUCT LIST

GEMCITABINE HYDROCHLORIDESOLUTION; INTRAVENOUS
INFUGEM

+	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

TABLET; ORAL

GEMFIBROZIL

AB	APOTEX	600MG	A075034 001	Jul 20, 1998
AB	AUROBINDO PHARMA LTD	600MG	A202726 001	Sep 16, 2015
AB	CADILA	600MG	A204189 001	Aug 28, 2018
AB	CADILA PHARMS LTD	600MG	A203266 001	Jun 17, 2016
AB	CARIBE HOLDINGS	600MG	A078012 001	Mar 26, 2007
AB	CHARTWELL MOLECULES	600MG	A074270 001	Sep 27, 1993
AB	HIKMA PHARMS	600MG	A078599 001	Aug 16, 2010
AB	IMPAX PHARMS	600MG	A078207 001	Jun 01, 2007
AB	INVAGEN PHARMS	600MG	A077836 001	Jul 27, 2006
AB	NORTHSTAR HLTHCARE	600MG	A079072 001	Sep 13, 2010

LOPID

AB	+	PFIZER PHARMS	600MG	N018422 003	Nov 20, 1986
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GEMIFLOXACIN MESYLATE

TABLET; ORAL

FACTIVE

AB	+	LG CHEM LTD	EQ 320MG BASE	N021158 001	Apr 04, 2003
AB		ORCHID HLTHCARE	EQ 320MG BASE	A090466 001	Jun 15, 2015

GENTAMICIN SULFATE

CREAM; TOPICAL

GENTAMICIN SULFATE

AT		ACP NIMBLE	EQ 0.1% BASE	A064056 001	Apr 29, 1994
AT	!	PERRIGO NEW YORK	EQ 0.1% BASE	A062307 001	
AT		TELIGENT PHARMA INC	EQ 0.1% BASE	A209304 001	Oct 18, 2019

INJECTABLE; INJECTION

GENTAMICIN SULFATE

AP	!	PRESENIUS KABI USA	EQ 10MG BASE/ML	A062366 002	Feb 06, 1986
AP	!		EQ 40MG BASE/ML	A062366 001	Aug 04, 1983
AP		HOSPIRA	EQ 10MG BASE/ML	A062420 001	Aug 15, 1983
AP			EQ 10MG BASE/ML	A062612 004	Feb 20, 1986
AP			EQ 40MG BASE/ML	A062420 002	Aug 15, 1983

GENTAMICIN SULFATE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

AP		BAXTER HLTHCARE	EQ 1.2MG BASE/ML	A062373 007	Sep 07, 1982
AP			EQ 1.6MG BASE/ML	A062373 008	Sep 07, 1982
AP			EQ 80MG BASE/100ML	A062373 002	Sep 07, 1982
AP			EQ 100MG BASE/100ML	A062373 005	Sep 07, 1982
AP		HOSPIRA	EQ 1.2MG BASE/ML	A062414 001	Aug 15, 1983
AP			EQ 1.6MG BASE/ML	A062414 003	Aug 15, 1983
AP			EQ 80MG BASE/100ML	A062414 008	Aug 15, 1983
AP			EQ 100MG BASE/100ML	A062414 010	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

AT	!	AKORN	EQ 0.3% BASE	A064093 001	Aug 31, 1995
AT		FERA PHARMS LLC	EQ 0.3% BASE	A065024 001	Jul 30, 2004

OINTMENT; TOPICAL

GENTAMICIN SULFATE

AT		ACP NIMBLE	EQ 0.1% BASE	A064054 001	Apr 29, 1994
AT		FOUGERA PHARMS INC	EQ 0.1% BASE	A062533 001	Oct 05, 1984
AT	!	PERRIGO NEW YORK	EQ 0.1% BASE	A062351 001	Feb 18, 1982
AT		TARO	EQ 0.1% BASE	A062477 001	Dec 23, 1983
AT		TELIGENT PHARMA INC	EQ 0.1% BASE	A209233 001	Dec 31, 2018

SOLUTION/DROPS; OPHTHALMIC

GENOPTIC

AT	!	ALLERGAN	EQ 0.3% BASE	A062452 001	Oct 10, 1984
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GENTAK

AT		AKORN	EQ 0.3% BASE	A064163 001	Oct 12, 2001
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GENTAMICIN SULFATE

AT		AKORN	EQ 0.3% BASE	A062635 001	Jan 08, 1987
AT		BAUSCH AND LOMB	EQ 0.3% BASE	A064048 001	May 11, 1994
AT		PERRIGO CO	EQ 0.3% BASE	A065121 001	Jan 30, 2004

PRESCRIPTION DRUG PRODUCT LIST

GENTAMICIN SULFATE

SOLUTION/DROPS;OPHTHALMIC

GENTAMICIN SULFATE

TENNESSEE

AT SANDOZ INC **EQ 0.3% BASE** **A062196 001**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

GILTERITINIB FUMARATE

TABLET;ORAL

XOSPATA

+! ASTELLAS EQ 40MG BASE N211349 001 Nov 28, 2018

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 INC EQ 25MG BASE N210656 001 Nov 21, 2018

+! EQ 100MG BASE N210656 002 Nov 21, 2018

GLATIRAMER ACETATE

INJECTABLE;SUBCUTANEOUS

COPAXONE**AP** +! TEVA PHARMS USA **20MG/ML** **N020622 002** Feb 12, 2002**AP** +! **40MG/ML** **N020622 003** Jan 28, 2014GLATIRAMER ACETATE**AP** MYLAN **20MG/ML** **A091646 001** Oct 03, 2017**AP** **40MG/ML** **A206936 001** Oct 03, 2017GLATOPIA**AP** SANDOZ INC **20MG/ML** **A090218 001** Apr 16, 2015**AP** **40MG/ML** **A206921 001** Feb 12, 2018GLECAPREVIR; PIBRENTASVIR

TABLET;ORAL

MAVYRET

+! ABBVIE INC 100MG;40MG N209394 001 Aug 03, 2017

GLIMEPIRIDE

TABLET;ORAL

AMARYL**AB** +! SANOFI AVENTIS US **1MG** **N020496 001** Nov 30, 1995**AB** + **2MG** **N020496 002** Nov 30, 1995**AB** + **4MG** **N020496 003** Nov 30, 1995GLIMEPIRIDE**AB** ACCORD HLTHCARE **1MG** **A078181 001** Aug 23, 2007**AB** **2MG** **A078181 002** Aug 23, 2007**AB** **4MG** **A078181 003** Aug 23, 2007**AB** AUROBINDO PHARMA LTD **1MG** **A202759 001** Jun 29, 2012**AB** **2MG** **A202759 002** Jun 29, 2012**AB** **4MG** **A202759 003** Jun 29, 2012**AB** CARLSBAD **1MG** **A077911 001** Sep 22, 2009**AB** **2MG** **A077911 002** Sep 22, 2009**AB** **4MG** **A077911 003** Sep 22, 2009**AB** DR REDDYS LABS LTD **1MG** **A077091 001** Oct 06, 2005**AB** **2MG** **A077091 002** Oct 06, 2005**AB** **4MG** **A077091 003** Oct 06, 2005**AB** INDOCO REMEDIES **1MG** **A202112 001** Apr 17, 2013**AB** **2MG** **A202112 002** Apr 17, 2013**AB** **4MG** **A202112 003** Apr 17, 2013**AB** INVAGEN PHARMS **1MG** **A077295 001** Oct 06, 2005**AB** **2MG** **A077295 002** Oct 06, 2005**AB** **4MG** **A077295 003** Oct 06, 2005**AB** MICRO LABS **1MG** **A091220 001** Jun 29, 2012**AB** **2MG** **A091220 002** Jun 29, 2012**AB** **4MG** **A091220 004** Jun 29, 2012

PRESCRIPTION DRUG PRODUCT LIST

GLIMEPIRIDE

TABLET; ORAL

GLIMEPIRIDE

<u>AB</u>		<u>8MG</u>	<u>A091220 006</u>	Jun 29, 2012
<u>AB</u>	MYLAN	<u>1MG</u>	<u>A077624 001</u>	Nov 28, 2005
<u>AB</u>		<u>2MG</u>	<u>A077624 002</u>	Nov 28, 2005
<u>AB</u>		<u>4MG</u>	<u>A077624 003</u>	Nov 28, 2005
<u>AB</u>	PRINSTON INC	<u>1MG</u>	<u>A077370 001</u>	Dec 23, 2005
<u>AB</u>		<u>2MG</u>	<u>A077370 002</u>	Dec 23, 2005
<u>AB</u>		<u>4MG</u>	<u>A077370 003</u>	Dec 23, 2005
<u>AB</u>		<u>8MG</u>	<u>A077370 004</u>	Dec 23, 2005
	MICRO LABS	3MG	A091220 003	Jun 29, 2012
		6MG	A091220 005	Jun 29, 2012

GLIMEPIRIDE; PIOGLITAZONE HYDROCHLORIDE

TABLET; ORAL

DUETACT

<u>AB</u>	+!	TAKEDA PHARMS USA	<u>2MG;30MG</u>	<u>N021925 001</u>	Jul 28, 2006
<u>AB</u>	+		<u>4MG;30MG</u>	<u>N021925 002</u>	Jul 28, 2006

PIOGLITAZONE HYDROCHLORIDE AND GLIMEPIRIDE

<u>AB</u>		SANDOZ	<u>2MG;30MG</u>	<u>A201049 001</u>	Jan 04, 2013
<u>AB</u>			<u>4MG;30MG</u>	<u>A201049 002</u>	Jan 04, 2013

GLIPIZIDE

TABLET; ORAL

GLIPIZIDE

<u>AB</u>		ACCORD HLTHCARE	<u>5MG</u>	<u>A074550 001</u>	Sep 11, 1997
<u>AB</u>			<u>10MG</u>	<u>A074550 002</u>	Sep 11, 1997
<u>AB</u>		APOTEX	<u>5MG</u>	<u>A075795 001</u>	Jun 13, 2001
<u>AB</u>			<u>10MG</u>	<u>A075795 002</u>	Jun 13, 2001
<u>AB</u>		MYLAN	<u>5MG</u>	<u>A074226 001</u>	May 10, 1994
<u>AB</u>			<u>10MG</u>	<u>A074226 002</u>	May 10, 1994
<u>AB</u>		SANDOZ	<u>5MG</u>	<u>A074305 001</u>	Apr 07, 1995
<u>AB</u>			<u>10MG</u>	<u>A074305 002</u>	Apr 07, 1995
<u>AB</u>		SUN PHARM INDS INC	<u>5MG</u>	<u>A077820 001</u>	Jul 11, 2006
<u>AB</u>			<u>10MG</u>	<u>A077820 002</u>	Jul 11, 2006
<u>AB</u>		WATSON LABS TEVA	<u>5MG</u>	<u>A074223 001</u>	Feb 27, 1995
<u>AB</u>			<u>10MG</u>	<u>A074223 002</u>	Feb 27, 1995

GLUCOTROL

<u>AB</u>	+	PFIZER	<u>5MG</u>	<u>N017783 001</u>	May 08, 1984
<u>AB</u>	+!		<u>10MG</u>	<u>N017783 002</u>	May 08, 1984

TABLET, EXTENDED RELEASE; ORAL

GLIPIZIDE

<u>AB</u>		AUROBINDO PHARMA LTD	<u>2.5MG</u>	<u>A206928 001</u>	May 12, 2017
<u>AB</u>			<u>5MG</u>	<u>A206928 002</u>	May 12, 2017
<u>AB</u>			<u>10MG</u>	<u>A206928 003</u>	May 12, 2017
<u>AB</u>		PAR PHARM	<u>5MG</u>	<u>A076159 002</u>	Sep 20, 2013
<u>AB</u>			<u>10MG</u>	<u>A076159 001</u>	Sep 20, 2013
<u>AB</u>		UNIQUE PHARM LABS	<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

GLUCOTROL XL

<u>AB</u>	+	PFIZER	<u>2.5MG</u>	<u>N020329 003</u>	Aug 10, 1999
<u>AB</u>	+		<u>5MG</u>	<u>N020329 001</u>	Apr 26, 1994
<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 PHARMS INC	<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

PRESCRIPTION DRUG PRODUCT LIST

GLIPIZIDE; METFORMIN HYDROCHLORIDE

TABLET; ORAL

GLIPIZIDE AND METFORMIN HYDROCHLORIDE

AB	!		<u>5MG;500MG</u>	<u>A077270 003</u>	Oct 28, 2005
AB		ZYDUS PHARMS USA INC	<u>2.5MG;250MG</u>	<u>A078905 001</u>	Jan 31, 2011
AB			<u>2.5MG;500MG</u>	<u>A078905 002</u>	Jan 31, 2011
AB			<u>5MG;500MG</u>	<u>A078905 003</u>	Jan 31, 2011

GLUCAGON

INJECTABLE; INJECTION

GLUCAGON

+! LILLY

1MG/VIAL

N020928 001 Sep 11, 1998

POWDER; NASAL

BAQSIMI

+! ELI LILLY AND CO

3MG

N210134 001 Jul 24, 2019

SOLUTION; SUBCUTANEOUS

GVOKE HYPOPEN

+! 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 PFS

+! XERIS

0.5MG/0.1ML (0.5MG/0.1ML)

N212097 001 Sep 10, 2019

+!

1MG/0.2ML (1MG/0.2ML)

N212097 002 Sep 10, 2019

GLUCAGON HYDROCHLORIDE

INJECTABLE; INJECTION

GLUCAGEN

+! NOVO NORDISK

EQ 1MG BASE/VIAL

N020918 001 Jun 22, 1998

POWDER; INTRAMUSCULAR, INTRAVENOUS

GLUCAGON

+! FRESENIUS KABI USA

EQ 1MG BASE/VIAL

N201849 001 May 08, 2015

GLYBURIDE

TABLET; ORAL

GLYBURIDE (MICRONIZED)

AB		DAVA PHARMS INC	<u>1.5MG</u>	<u>A074591 001</u>	Dec 22, 1997
AB			<u>3MG</u>	<u>A074591 002</u>	Dec 22, 1997
AB			<u>4.5MG</u>	<u>A074591 003</u>	Dec 22, 1997
AB			<u>6MG</u>	<u>A074591 004</u>	Dec 22, 1997
AB		HIKMA	<u>1.5MG</u>	<u>A075890 001</u>	Jul 31, 2003
AB			<u>3MG</u>	<u>A075890 002</u>	Jul 31, 2003
AB			<u>6MG</u>	<u>A075890 003</u>	Jul 31, 2003
AB		TEVA	<u>1.5MG</u>	<u>A074686 001</u>	Apr 20, 1999
AB			<u>3MG</u>	<u>A074686 002</u>	Apr 20, 1999
AB			<u>4.5MG</u>	<u>A074686 003</u>	Apr 20, 1999
AB			<u>6MG</u>	<u>A074686 004</u>	Apr 20, 1999

GLYNASE

AB	+	PHARMACIA AND UPJOHN	<u>1.5MG</u>	<u>N020051 001</u>	Mar 04, 1992
AB	+		<u>3MG</u>	<u>N020051 002</u>	Mar 04, 1992
AB	+!		<u>6MG</u>	<u>N020051 004</u>	Sep 24, 1993

GLYBURIDE

AB1		AUROBINDO PHARMA	<u>1.25MG</u>	<u>A077537 001</u>	Oct 18, 2007
AB1			<u>2.5MG</u>	<u>A077537 002</u>	Oct 18, 2007
AB1			<u>5MG</u>	<u>A077537 003</u>	Oct 18, 2007
AB1		CADILA PHARMS LTD	<u>1.25MG</u>	<u>A203379 001</u>	Jan 04, 2019
AB1			<u>2.5MG</u>	<u>A203379 002</u>	Jan 04, 2019
AB1			<u>5MG</u>	<u>A203379 003</u>	Jan 04, 2019
AB1		EPIC PHARMA LLC	<u>1.25MG</u>	<u>A076257 001</u>	Jun 27, 2002
AB1			<u>2.5MG</u>	<u>A076257 002</u>	Jun 27, 2002
AB1			<u>5MG</u>	<u>A076257 003</u>	Jun 27, 2002
AB1		HERITAGE PHARMS INC	<u>1.25MG</u>	<u>A090937 001</u>	Feb 28, 2011
AB1			<u>2.5MG</u>	<u>A090937 002</u>	Feb 28, 2011
AB1			<u>5MG</u>	<u>A090937 003</u>	Feb 28, 2011
AB1		PHARMADAX INC	<u>1.25MG</u>	<u>A203581 001</u>	Apr 14, 2016
AB1			<u>2.5MG</u>	<u>A203581 002</u>	Apr 14, 2016
AB1			<u>5MG</u>	<u>A203581 003</u>	Apr 14, 2016
AB1		TEVA	<u>1.25MG</u>	<u>A074388 001</u>	Aug 29, 1995
AB1			<u>2.5MG</u>	<u>A074388 002</u>	Aug 29, 1995
AB1	!		<u>5MG</u>	<u>A074388 003</u>	Aug 29, 1995
AB1		ZYDUS PHARMS	<u>1.25mg</u>	<u>A206749 001</u>	May 10, 2016
AB1			<u>2.5mg</u>	<u>A206749 002</u>	May 10, 2016
AB1			<u>5MG</u>	<u>A206749 003</u>	May 10, 2016

PRESCRIPTION DRUG PRODUCT LIST

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GLYBURIDE

TABLET; ORAL

DIABETA

AB2	+	SANOFI AVENTIS US	<u>1.25MG</u>	<u>N017532</u>	<u>001</u>	May 01, 1984
AB2	+		<u>2.5MG</u>	<u>N017532</u>	<u>002</u>	May 01, 1984
AB2	+	!	<u>5MG</u>	<u>N017532</u>	<u>003</u>	May 01, 1984

GLYBURIDE

AB2		IMPAX LABS INC	<u>1.25MG</u>	<u>A206079</u>	<u>001</u>	Sep 30, 2015
AB2			<u>2.5MG</u>	<u>A206079</u>	<u>002</u>	Sep 30, 2015
AB2			<u>5MG</u>	<u>A206079</u>	<u>003</u>	Sep 30, 2015

GLYBURIDE; METFORMIN HYDROCHLORIDE

TABLET; ORAL

GLYBURIDE AND METFORMIN HYDROCHLORIDE

AB		ACTAVIS ELIZABETH	<u>1.25MG;250MG</u>	<u>A076716</u>	<u>001</u>	Jun 28, 2005
AB			<u>2.5MG;500MG</u>	<u>A076716</u>	<u>002</u>	Jun 28, 2005
AB			<u>5MG;500MG</u>	<u>A076716</u>	<u>003</u>	Jun 28, 2005
AB		AUROBINDO PHARMA	<u>1.25MG;250MG</u>	<u>A077870</u>	<u>001</u>	Nov 14, 2007
AB		!	<u>2.5MG;500MG</u>	<u>A077870</u>	<u>002</u>	Nov 14, 2007
AB			<u>5MG;500MG</u>	<u>A077870</u>	<u>003</u>	Nov 14, 2007
AB		IMPAX LABS INC	<u>1.25MG;250MG</u>	<u>A076345</u>	<u>001</u>	Feb 18, 2004
AB			<u>2.5MG;500MG</u>	<u>A076345</u>	<u>002</u>	Feb 18, 2004
AB			<u>5MG;500MG</u>	<u>A076345</u>	<u>003</u>	Feb 18, 2004
AB		ZYDUS PHARMS	<u>1.25MG;250MG</u>	<u>A206748</u>	<u>001</u>	Feb 29, 2016
AB			<u>2.5MG;500MG</u>	<u>A206748</u>	<u>002</u>	Feb 29, 2016
AB			<u>5MG;500MG</u>	<u>A206748</u>	<u>003</u>	Feb 29, 2016

GLYCEROL PHENYLBUTYRATE

LIQUID; ORAL

RAVICTI

+	!	HORIZON THERAP	1.1GM/ML	N203284	001	Feb 01, 2013
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GLYCINE

SOLUTION; IRRIGATION

AMINOACETIC ACID 1.5% IN PLASTIC CONTAINER

AT	+	!	BAXTER HLTHCARE	<u>1.5GM/100ML</u>	<u>N017865</u>	<u>001</u>
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GLYCINE 1.5% IN PLASTIC CONTAINER

AT			B BRAUN	<u>1.5GM/100ML</u>	<u>N016784</u>	<u>001</u>
AT			ICU MEDICAL INC	<u>1.5GM/100ML</u>	<u>N018315</u>	<u>001</u>

GLYCOPYRROLATE

INJECTABLE; INJECTION

GLYCOPYRROLATE

AP			AM REGENT	<u>0.2MG/ML</u>	<u>A089335</u>	<u>001</u>	Jul 23, 1986
AP			AMNEAL	<u>0.2MG/ML</u>	<u>A208973</u>	<u>001</u>	Jun 15, 2017
AP			APOTEX	<u>0.2MG/ML</u>	<u>A210246</u>	<u>001</u>	Oct 29, 2019
AP			AUROBINDO PHARMA LTD	<u>0.2MG/ML</u>	<u>A210244</u>	<u>001</u>	Nov 28, 2018
AP			CAPLIN	<u>0.2MG/ML</u>	<u>A211705</u>	<u>001</u>	Mar 20, 2019
AP			FRESENIUS KABI USA	<u>0.2MG/ML</u>	<u>A209024</u>	<u>001</u>	Oct 31, 2018
AP				<u>0.2MG/ML</u>	<u>A209328</u>	<u>001</u>	Oct 27, 2017
AP			GLAND PHARMA LTD	<u>0.2MG/ML</u>	<u>A212612</u>	<u>001</u>	Sep 30, 2019
AP		!	HIKMA FARMACEUTICA	<u>0.2MG/ML</u>	<u>A090963</u>	<u>001</u>	Sep 21, 2011
AP			PIRAMAL CRITICAL	<u>0.2MG/ML</u>	<u>A210842</u>	<u>001</u>	Oct 25, 2018
AP			PRINSTON INC	<u>0.2MG/ML</u>	<u>A210927</u>	<u>001</u>	Oct 31, 2018
AP			RICONPHARMA LLC	<u>0.2MG/ML</u>	<u>A210083</u>	<u>001</u>	Feb 21, 2020
AP			SANDOZ INC	<u>0.2MG/ML</u>	<u>A211334</u>	<u>001</u>	May 14, 2019
AP			SOMERSET THERAPS LLC	<u>0.2MG/ML</u>	<u>A207639</u>	<u>001</u>	Jun 23, 2017

POWDER; INHALATION

SEEBRI

+	!	SUNOVION PHARMS INC	15.6MCG/INH	N207923	001	Oct 29, 2015
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SOLUTION; INHALATION

LONHALA MAGNAIR KIT

+	!	SUNOVION RESP	25MCG/ML	N208437	001	Dec 05, 2017
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SOLUTION; INTRAMUSCULAR, INTRAVENOUS

GLYRX-PF

		EXELA PHARMA SCS LLC	0.2MG/ML (0.2MG/ML)	N210997	001	Jul 11, 2018
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			0.4MG/2ML (0.2MG/ML)	N210997	002	Jul 11, 2018
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SOLUTION; ORAL

CUVPOSA

+	!	MERZ PHARMS	1MG/5ML	N022571	001	Jul 28, 2010
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PRESCRIPTION DRUG PRODUCT LISTGLYCOPYRROLATE

TABLET; ORAL

GLYCOPYRROLATE

<u>AA</u>	AUROLIFE PHARMA LLC	<u>1MG</u>	<u>A202675 001</u>	Apr 15, 2013
<u>AA</u>		<u>2MG</u>	<u>A202675 002</u>	Oct 30, 2018
<u>AA</u>	DR REDDYS LABS LTD	<u>1MG</u>	<u>A040847 001</u>	Mar 21, 2008
<u>AA</u>		<u>2MG</u>	<u>A040847 002</u>	Mar 21, 2008
<u>AA</u>	HERITAGE PHARMS INC	<u>1MG</u>	<u>A207201 001</u>	Jan 03, 2017
<u>AA</u>		<u>2MG</u>	<u>A207201 002</u>	Jan 03, 2017
<u>AA</u>	KENTON	<u>1MG</u>	<u>A091182 001</u>	Feb 03, 2014
<u>AA</u>		<u>2MG</u>	<u>A091182 002</u>	Feb 03, 2014
<u>AA</u>	LEADING PHARMA LLC	<u>1MG</u>	<u>A090195 001</u>	Sep 21, 2012
<u>AA</u>		<u>2MG</u>	<u>A090195 002</u>	Sep 21, 2012
<u>AA</u>	NATCO PHARMA LTD	<u>1MG</u>	<u>A091413 001</u>	Jun 20, 2016
<u>AA</u>		<u>2MG</u>	<u>A091413 002</u>	Jun 20, 2016
<u>AA</u>	ORIT LABS LLC	<u>1MG</u>	<u>A203657 001</u>	Nov 30, 2018
<u>AA</u>		<u>2MG</u>	<u>A203657 002</u>	Nov 30, 2018
<u>AA</u>	OXFORD PHARMS	<u>1MG</u>	<u>A090020 001</u>	Oct 19, 2011
<u>AA</u>		<u>2MG</u>	<u>A090020 002</u>	Oct 19, 2011
<u>AA</u>	! PAR PHARM	<u>1MG</u>	<u>A040653 001</u>	Aug 31, 2006
<u>AA</u>	!	<u>2MG</u>	<u>A040653 002</u>	Aug 31, 2006
<u>AA</u>	RISING	<u>1MG</u>	<u>A040821 001</u>	Dec 29, 2008
<u>AA</u>		<u>2MG</u>	<u>A040821 002</u>	Dec 29, 2008
<u>AA</u>	SUN PHARM INDS LTD	<u>1MG</u>	<u>A040844 001</u>	Aug 18, 2009
<u>AA</u>		<u>2MG</u>	<u>A040844 002</u>	Aug 18, 2009
	NEXGEN PHARMA	1.5MG	A091522 001	Mar 12, 2012

GLYCOPYRROLATE; INDACATEROL MALEATE

POWDER; INHALATION

UTIBRON

+! SUNOVION PHARMS INC 15.6MCG/INH;27.5MCG/INH N207930 001 Oct 29, 2015

GLYCOPYRRONIUM TOSYLATE

CLOTH; TOPICAL

QBREXZA

+! DERMIRA INC 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

GONADOTROPIN, CHORIONIC

INJECTABLE; INJECTION

CHORIONIC GONADOTROPIN

<u>AP</u>	+! FERRING	<u>10,000 UNITS/VIAL</u>	<u>N017016 007</u>	
<u>AP</u>	+! FRESENIUS KABI USA	<u>10,000 UNITS/VIAL</u>	<u>N017067 002</u>	
	<u>PREGNYL</u>			
<u>AP</u>	+! ORGANON USA INC	<u>10,000 UNITS/VIAL</u>	<u>N017692 001</u>	
	CHORIONIC GONADOTROPIN			
	+! FERRING	5,000 UNITS/VIAL	N017016 006	

GOSERELIN ACETATE

IMPLANT; IMPLANTATION

ZOLADEX

+! TERSERA THERAPS LLC EQ 3.6MG BASE N019726 001 Dec 29, 1989
+! EQ 10.8MG BASE N020578 001 Jan 11, 1996GRAMICIDIN; NEOMYCIN SULFATE; POLYMYXIN B SULFATE

SOLUTION/DROPS; OPHTHALMIC

NEOMYCIN AND POLYMYXIN B SULFATES AND GRAMICIDIN

<u>AT</u>	AMRING PHARMS	<u>0.025MG/ML;EQ 1.75MG BASE/ML;10,000 UNITS/ML</u>	<u>A065187 001</u>	Oct 28, 2005
<u>AT</u>	! BAUSCH AND LOMB	<u>0.025MG/ML;EQ 1.75MG BASE/ML;10,000 UNITS/ML</u>	<u>A064047 001</u>	Jan 31, 1996
	<u>NEOSPORIN</u>			
<u>AT</u>	! MONARCH PHARMS	<u>0.025MG/ML;EQ 1.75MG BASE/ML;10,000 UNITS/ML</u>	<u>A060582 001</u>	

GRANISETRON

FILM, EXTENDED RELEASE; TRANSDERMAL

SANCUSO

+! KYOWA KIRIN 3.1MG/24HR N022198 001 Sep 12, 2008

INJECTION, EXTENDED RELEASE; SUBCUTANEOUS

SUSTOL

+! HERON THERAPS INC 10MG/0.4ML (10MG/0.4ML) N022445 001 Aug 09, 2016

PRESCRIPTION DRUG PRODUCT LIST

3-215 (of 453)

GRANISETRON HYDROCHLORIDE

INJECTABLE; INJECTION

GRANISETRON HYDROCHLORIDE

AP	AKORN INC	<u>EQ 0.1MG BASE/ML (EQ 0.1MG BASE/ML)</u>	<u>A079119 001</u>	Sep 10, 2009
AP		<u>EQ 1MG BASE/ML (EQ 1MG BASE/ML)</u>	<u>A079078 001</u>	Sep 14, 2009
AP		<u>EQ 4MG BASE/4ML (EQ 1MG BASE/ML)</u>	<u>A079078 002</u>	Sep 14, 2009
AP	AUROBINDO PHARMA LTD	<u>EQ 1MG BASE/ML (EQ 1MG BASE/ML)</u>	<u>A204238 001</u>	Jul 06, 2016
AP		<u>EQ 4MG BASE/4ML (EQ 1MG BASE/ML)</u>	<u>A204238 002</u>	Jul 06, 2016
AP	BIONPHARMA INC	<u>EQ 0.1MG BASE/ML (EQ 0.1MG BASE/ML)</u>	<u>A078863 001</u>	Jun 30, 2008
AP		<u>EQ 4MG BASE/4ML (EQ 1MG BASE/ML)</u>	<u>A078880 001</u>	Jun 30, 2008
AP	CIPLA LTD	<u>EQ 0.1MG BASE/ML (EQ 0.1MG BASE/ML)</u>	<u>A078262 001</u>	Dec 31, 2007
AP		<u>EQ 1MG BASE/ML (EQ 1MG BASE/ML)</u>	<u>A078258 001</u>	Jun 30, 2008
AP		<u>EQ 4MG BASE/4ML (EQ 1MG BASE/ML)</u>	<u>A078258 002</u>	Jun 30, 2008
AP	! DR REDDYS	<u>EQ 0.1MG BASE/ML (EQ 0.1MG BASE/ML)</u>	<u>A078392 001</u>	Dec 31, 2007
AP	!	<u>EQ 4MG BASE/4ML (EQ 1MG BASE/ML)</u>	<u>A077297 001</u>	Jun 30, 2008
AP	FRESENIUS KABI USA	<u>EQ 0.1MG BASE/ML (EQ 0.1MG BASE/ML)</u>	<u>A078522 001</u>	Dec 31, 2007
AP		<u>EQ 4MG BASE/4ML (EQ 1MG BASE/ML)</u>	<u>A078090 001</u>	Jun 30, 2008
AP	HIKMA FARMACEUTICA	<u>EQ 1MG BASE/ML (EQ 1MG BASE/ML)</u>	<u>A078629 001</u>	Dec 23, 2009
AP		<u>EQ 4MG BASE/4ML (EQ 1MG BASE/ML)</u>	<u>A078629 002</u>	Dec 23, 2009
AP	MYLAN ASI	<u>EQ 0.1MG BASE/ML (EQ 0.1MG BASE/ML)</u>	<u>A091136 001</u>	Apr 09, 2010
AP		<u>EQ 1MG BASE/ML (EQ 1MG BASE/ML)</u>	<u>A091136 002</u>	Apr 09, 2010
AP		<u>EQ 4MG BASE/4ML (EQ 1MG BASE/ML)</u>	<u>A091137 002</u>	Apr 09, 2010
AP	MYLAN LABS LTD	<u>EQ 0.1MG BASE/ML (EQ 0.1MG BASE/ML)</u>	<u>A203454 001</u>	Apr 04, 2017
AP		<u>EQ 1MG BASE/ML (EQ 1MG BASE/ML)</u>	<u>A203454 002</u>	Apr 04, 2017
AP		<u>EQ 4MG BASE/4ML (EQ 1MG BASE/ML)</u>	<u>A203453 001</u>	Jan 31, 2017
AP	SANDOZ INC	<u>EQ 0.1MG BASE/ML (EQ 0.1MG BASE/ML)</u>	<u>A078534 001</u>	Apr 30, 2009
AP		<u>EQ 1MG BASE/ML (EQ 1MG BASE/ML)</u>	<u>A078531 001</u>	Apr 30, 2009
AP		<u>EQ 1MG BASE/ML (EQ 1MG BASE/ML)</u>	<u>A078835 001</u>	Jun 30, 2008
AP		<u>EQ 4MG BASE/4ML (EQ 1MG BASE/ML)</u>	<u>A078531 002</u>	Apr 30, 2009
AP		<u>EQ 4MG BASE/4ML (EQ 1MG BASE/ML)</u>	<u>A078835 002</u>	Jun 30, 2008
AP	WEST-WARD PHARMS INT	<u>EQ 0.1MG BASE/ML (EQ 0.1MG BASE/ML)</u>	<u>A077913 001</u>	Jun 26, 2008
AP		<u>EQ 4MG BASE/4ML (EQ 1MG BASE/ML)</u>	<u>A077186 001</u>	Jun 30, 2008
AP		<u>EQ 1MG BASE/ML (EQ 1MG BASE/ML)</u>	<u>A077187 001</u>	Jun 30, 2008
AP		<u>EQ 4MG BASE/4ML (EQ 1MG BASE/ML)</u>	<u>A077177 001</u>	Dec 31, 2007
AP	WOCKHARDT USA	<u>EQ 0.1MG BASE/ML (EQ 0.1MG BASE/ML)</u>	<u>A078566 001</u>	Feb 29, 2008
AP		<u>EQ 1MG BASE/ML (EQ 1MG BASE/ML)</u>	<u>A078564 001</u>	Jun 30, 2008
AP		<u>EQ 4MG BASE/4ML (EQ 1MG BASE/ML)</u>	<u>A078565 001</u>	Jun 30, 2008
AP	YUNG SHIN PHARM	<u>EQ 0.1MG BASE/ML (EQ 0.1MG BASE/ML)</u>	<u>A202647 001</u>	Mar 06, 2020

GRANISETRON HYDROCHLORIDE PRESERVATIVE FREE

AP	BIONPHARMA INC	<u>EQ 1MG BASE/ML (EQ 1MG BASE/ML)</u>	<u>A078863 002</u>	Jun 30, 2008
AP	! FRESENIUS KABI USA	<u>EQ 1MG BASE/ML (EQ 1MG BASE/ML)</u>	<u>A078096 001</u>	Jun 30, 2008

TABLET; ORAL

GRANISETRON HYDROCHLORIDE

AB	APOTEX INC	<u>EQ 1MG BASE</u>	<u>A078843 001</u>	Feb 27, 2008
AB	CHARTWELL MOLECULAR	<u>EQ 1MG BASE</u>	<u>A078037 001</u>	Feb 27, 2008
AB	DR REDDYS LABS LTD	<u>EQ 1MG BASE</u>	<u>A078846 001</u>	Feb 27, 2009
AB	! HIKMA	<u>EQ 1MG BASE</u>	<u>A077842 001</u>	Dec 31, 2007
AB	NATCO PHARMA	<u>EQ 1MG BASE</u>	<u>A078969 001</u>	Jun 22, 2009
AB	ORCHID HLTHCARE	<u>EQ 1MG BASE</u>	<u>A078678 001</u>	Feb 13, 2008
AB	TARO	<u>EQ 1MG BASE</u>	<u>A090817 001</u>	May 28, 2010

GRISEOFULVIN, MICROSIZE

SUSPENSION; ORAL

GRISEOFULVIN

AB	! ACTAVIS MID ATLANTIC	<u>125MG/5ML</u>	<u>A065394 001</u>	Jul 06, 2007
AB	CHARTWELL RX	<u>125MG/5ML</u>	<u>A065200 001</u>	Mar 02, 2005
AB	CIPLA	<u>125MG/5ML</u>	<u>A065354 001</u>	Sep 10, 2007
AB	VINTAGE PHARMS	<u>125MG/5ML</u>	<u>A065438 001</u>	Oct 08, 2010

TABLET; ORAL

GRISEOFULVIN

AB	! SANDOZ INC	<u>500MG</u>	<u>A091592 002</u>	Aug 07, 2013
AB	SIGMAPHARM LABS LLC	<u>500MG</u>	<u>A202482 001</u>	Oct 22, 2012
	SANDOZ INC	<u>250MG</u>	<u>A091592 001</u>	Aug 07, 2013

GRISEOFULVIN, ULTRAMICROSIZE

TABLET; ORAL

GRIS-PEG

AB	+ VALEANT PHARMS INC	<u>125MG</u>	<u>N050475 001</u>	
AB	+!	<u>250MG</u>	<u>N050475 002</u>	

PRESCRIPTION DRUG PRODUCT LIST

3-216 (of 453)

GRISEOFULVIN, ULTRAMICROSIZED

TABLET; ORAL

GRISEOFULVIN, ULTRAMICROSIZED

<u>AB</u>	MOUNTAIN	<u>125MG</u>	<u>A204371 001</u>	Jan 09, 2014
<u>AB</u>		<u>250MG</u>	<u>A204371 002</u>	Jan 09, 2014
<u>AB</u>	SANDOZ INC	<u>125MG</u>	<u>A202805 001</u>	Dec 26, 2018
<u>AB</u>		<u>250MG</u>	<u>A202805 002</u>	Dec 26, 2018
<u>GRISEOFULVIN, ULTRAMICROSIZED</u>				
<u>AB</u>	SIGMAPHARM LABS LLC	<u>125MG</u>	<u>A202545 001</u>	Oct 22, 2012
<u>AB</u>		<u>250MG</u>	<u>A202545 002</u>	Oct 22, 2012

GUANFACINE HYDROCHLORIDE

TABLET; ORAL

GUANFACINE HYDROCHLORIDE

<u>AB</u>	AMNEAL PHARM	<u>EQ 1MG BASE</u>	<u>A075109 001</u>	Nov 25, 1998
<u>AB</u>		<u>EQ 2MG BASE</u>	<u>A075109 002</u>	Nov 25, 1998
<u>AB</u>	EPIC PHARMA LLC	<u>EQ 1MG BASE</u>	<u>A074673 001</u>	Feb 28, 1997
<u>AB</u>		<u>EQ 2MG BASE</u>	<u>A074673 002</u>	Feb 28, 1997
<u>AB</u>	MYLAN	<u>EQ 1MG BASE</u>	<u>A074796 001</u>	Jan 27, 1997
<u>AB</u>	!	<u>EQ 2MG BASE</u>	<u>A074796 002</u>	Jan 27, 1997
<u>AB</u>	WATSON LABS	<u>EQ 1MG BASE</u>	<u>A074145 001</u>	Oct 17, 1995
<u>AB</u>		<u>EQ 2MG BASE</u>	<u>A074145 002</u>	Oct 17, 1995

TABLET, EXTENDED RELEASE; ORAL

GUANFACINE HYDROCHLORIDE

<u>AB</u>	ACTAVIS ELIZABETH	<u>EQ 1MG BASE</u>	<u>A200881 001</u>	Oct 05, 2012
<u>AB</u>		<u>EQ 2MG BASE</u>	<u>A200881 002</u>	Oct 05, 2012
<u>AB</u>		<u>EQ 3MG BASE</u>	<u>A200881 003</u>	Oct 05, 2012
<u>AB</u>		<u>EQ 4MG BASE</u>	<u>A200881 004</u>	Oct 05, 2012
<u>AB</u>	APOTEX	<u>EQ 1MG BASE</u>	<u>A205430 001</u>	Jul 25, 2018
<u>AB</u>		<u>EQ 2MG BASE</u>	<u>A205430 002</u>	Jul 25, 2018
<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 INC	<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>INTUNIV</u>				
<u>AB</u>	+ SHIRE	<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

GUANIDINE HYDROCHLORIDE

TABLET; ORAL

GUANIDINE HYDROCHLORIDE

MERCK SHARP DOHME	125MG	N001546 001
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HALCINONIDE

CREAM; TOPICAL

HALCINONIDE

<u>AB</u>	MYLAN	<u>0.1%</u>	<u>A211027 001</u>	Aug 12, 2019
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HALOG

<u>AB</u>	+!	SUN PHARM INDS INC	<u>0.1%</u>	<u>N017556 001</u>
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OINTMENT; TOPICAL

HALOG

+!	SUN PHARM INDS INC	0.1%	N017824 001
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PRESCRIPTION DRUG PRODUCT LIST

HALOBETASOL PROPIONATE

AEROSOL, FOAM; TOPICAL

LEXETTE

+! MAYNE PHARMA 0.05% N210566 001 May 24, 2018

CREAM; TOPICAL

HALOBETASOL PROPIONATE**AB** ! ACP NIMBLE 0.05% **A078162 001** Apr 24, 2007**AB** FOUGERA PHARMS 0.05% **A077001 001** Dec 16, 2004**AB** PERRIGO ISRAEL 0.05% **A077123 001** Dec 16, 2004**AB** TARO 0.05% **A077227 001** Aug 04, 2005

LOTION; TOPICAL

BRYHALI

+! BAUSCH 0.01% N209355 001 Nov 06, 2018

ULTRAVATE

+! SUN PHARM 0.05% N208183 001 Nov 06, 2015

INDUSTRIES

OINTMENT; TOPICAL

HALOBETASOL PROPIONATE**AB** ACP NIMBLE 0.05% **A077109 001** Jun 14, 2005**AB** ! 0.05% **A077721 001** Sep 07, 2006**AB** PERRIGO ISRAEL 0.05% **A076872 001** Dec 16, 2004**AB** TARO 0.05% **A076994 001** Dec 16, 2004**AB** TELIGENT PHARMA INC 0.05% **A209978 001** Mar 20, 2018ULTRAVATE**AB** + SUN PHARM INDS INC 0.05% **N019968 001** Dec 17, 1990HALOBETASOL PROPIONATE; TAZAROTENE

LOTION; TOPICAL

DUOBRII

+! BAUSCH 0.01%; 0.045% N209354 001 Apr 25, 2019

HALOPERIDOL

TABLET; ORAL

HALOPERIDOL**AB** APPCO 0.5MG **A211061 001** Jan 08, 2020**AB** 1MG **A211061 002** Jan 08, 2020**AB** 2MG **A211061 003** Jan 08, 2020**AB** 5MG **A211061 004** Jan 08, 2020**AB** 10MG **A211061 005** Jan 08, 2020**AB** 20MG **A211061 006** Jan 08, 2020**AB** INNOGENIX 0.5MG **A071173 002** Jan 02, 1987**AB** 1MG **A071173 003** Jan 02, 1987**AB** 2MG **A071173 004** Jan 02, 1987**AB** 5MG **A071173 005** Jan 07, 1988**AB** 10MG **A071173 001** Jan 07, 1988**AB** 20MG **A071173 006** Jan 07, 1988**AB** MYLAN 0.5MG **A070278 006** Jun 10, 1986**AB** 1MG **A070278 004** Jun 10, 1986**AB** ! 2MG **A070278 001** Jun 10, 1986**AB** 5MG **A070278 005** Jun 10, 1986**AB** 10MG **A070278 002** Jul 16, 2009**AB** 20MG **A070278 003** Jul 16, 2009**AB** SANDOZ 0.5MG **A071206 001** Nov 17, 1986**AB** 1MG **A071207 001** Nov 17, 1986**AB** 5MG **A071209 001** Nov 17, 1986**AB** 10MG **A071210 001** Mar 11, 1988**AB** 20MG **A071211 001** Mar 11, 1988**AB** ZYDUS PHARMS USA 5MG **A077580 003** Nov 29, 2007**AB** 10MG **A077580 004** Nov 29, 2007**AB** 20MG **A077580 005** Nov 29, 2007HALOPERIDOL DECANOATE

INJECTABLE; INJECTION

HALDOL**AO** +! JANSSEN PHARMS EQ 50MG BASE/ML **N018701 001** Jan 14, 1986**AO** +! EQ 100MG BASE/ML **N018701 002** Jan 31, 1997HALOPERIDOL DECANOATE**AO** FRESENIUS KABI USA EQ 50MG BASE/ML **A074893 001** Dec 19, 1997**AO** EQ 100MG BASE/ML **A074893 002** Dec 19, 1997**AO** GLAND PHARMA LTD EQ 50MG BASE/ML **A205241 001** May 12, 2017**AO** EQ 100MG BASE/ML **A205241 002** May 12, 2017**AO** MYLAN LABS LTD EQ 50MG BASE/ML **A075440 001** Feb 28, 2000**AO** EQ 100MG BASE/ML **A075440 002** Feb 28, 2000**AO** SOMERSET THERAPS EQ 50MG BASE/ML **A209101 001** Jul 03, 2018

PRESCRIPTION DRUG PRODUCT LIST

HALOPERIDOL DECANOATE

INJECTABLE; INJECTION

HALOPERIDOL DECANOATE

LLC

<u>AO</u>		<u>EQ 100MG BASE/ML</u>	<u>A209101 002</u>	Jul 03, 2018
<u>AO</u>	TEVA PHARMS USA	<u>EQ 50MG BASE/ML</u>	<u>A075393 001</u>	May 11, 1999
<u>AO</u>		<u>EQ 100MG BASE/ML</u>	<u>A075393 002</u>	May 11, 1999
<u>AO</u>	WEST-WARD PHARMS INT	<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>	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

INJECTABLE; INJECTION

HALDOL

<u>AP</u>	+! JANSSEN PHARMS	<u>EQ 5MG BASE/ML</u>	<u>N015923 001</u>	
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HALOPERIDOL

<u>AP</u>	AKORN	<u>EQ 5MG BASE/ML</u>	<u>A204849 001</u>	Sep 06, 2017
<u>AP</u>	FRESENIUS KABI USA	<u>EQ 5MG BASE/ML</u>	<u>A075689 001</u>	Mar 09, 2001
<u>AP</u>	GLAND PHARMA LTD	<u>EQ 5MG BASE/ML</u>	<u>A076774 001</u>	Aug 25, 2004
<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
<u>AP</u>	TEVA PHARMS USA	<u>EQ 5MG BASE/ML</u>	<u>A076035 001</u>	Aug 29, 2001
<u>AP</u>	WEST-WARD PHARMS INT	<u>EQ 5MG BASE/ML</u>	<u>A075858 001</u>	Jun 18, 2001

HEPARIN SODIUM

INJECTABLE; INJECTION

HEPARIN SODIUM

<u>AP</u>	CASI PHARMS INC	<u>5,000 UNITS/ML</u>	<u>A091659 001</u>	Jun 08, 2011
<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 PHARMA LTD	<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>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>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
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HEPARIN SODIUM 1,000 UNITS IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

<u>AP</u>	+! B BRAUN	<u>200 UNITS/100ML</u>	<u>N019953 001</u>	Jul 20, 1992
<u>AP</u>	+! HOSPIRA	<u>200 UNITS/100ML</u>	<u>N018916 010</u>	Jun 23, 1989

PRESCRIPTION DRUG PRODUCT LIST

HEPARIN SODIUM

INJECTABLE; INJECTION

HEPARIN SODIUM 10,000 UNITS IN DEXTROSE 5% IN PLASTIC CONTAINER

<u>AP</u>	HOSPIRA	<u>10,000 UNITS/100ML</u>	<u>N019339 003</u>	Mar 27, 1985
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HEPARIN SODIUM 2,000 UNITS AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

<u>AP</u>	BAXTER HLTHCARE	<u>200 UNITS/100ML</u>	<u>N018609 002</u>	Apr 28, 1982
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HEPARIN SODIUM 2,000 UNITS IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

<u>AP</u>	+! HOSPIRA	<u>200 UNITS/100ML</u>	<u>N018916 011</u>	Jun 23, 1989
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HEPARIN SODIUM 20,000 UNITS IN DEXTROSE 5% IN PLASTIC CONTAINER

<u>AP</u>	+! B BRAUN	<u>4,000 UNITS/100ML</u>	<u>N019952 001</u>	Jul 20, 1992
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<u>AP</u>	HOSPIRA	<u>4,000 UNITS/100ML</u>	<u>N019805 001</u>	Jan 25, 1989
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HEPARIN SODIUM 25,000 UNITS IN DEXTROSE 5% IN PLASTIC CONTAINER

<u>AP</u>	+! B BRAUN	<u>5,000 UNITS/100ML</u>	<u>N019952 004</u>	Jul 20, 1992
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<u>AP</u>	+!	<u>10,000 UNITS/100ML</u>	<u>N019952 005</u>	Jul 20, 1992
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<u>AP</u>	HOSPIRA	<u>5,000 UNITS/100ML</u>	<u>N019339 004</u>	Mar 27, 1985
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<u>AP</u>		<u>5,000 UNITS/100ML</u>	<u>N019805 002</u>	Jan 25, 1989
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<u>AP</u>		<u>10,000 UNITS/100ML</u>	<u>N019339 002</u>	Mar 27, 1985
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HEPARIN SODIUM IN PLASTIC CONTAINER

<u>AP</u>	+! FRESENIUS KABI USA	<u>1,000 UNITS/ML</u>	<u>N017029 013</u>	Dec 05, 1985
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<u>AP</u>	+!	<u>5,000 UNITS/ML</u>	<u>N017029 014</u>	Dec 05, 1985
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<u>AP</u>	+!	<u>10,000 UNITS/ML</u>	<u>N017029 015</u>	Dec 05, 1985
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<u>AP</u>	+!	<u>20,000 UNITS/ML</u>	<u>N017029 016</u>	Dec 05, 1985
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HEPARIN SODIUM PRESERVATIVE FREE

<u>AP</u>	+! FRESENIUS KABI USA	<u>1,000 UNITS/ML</u>	<u>N017029 010</u>	Apr 28, 1986
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<u>AP</u>	SAGENT PHARMS	<u>1,000 UNITS/ML</u>	<u>A090810 001</u>	Jun 30, 2010
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<u>AP</u>	SHENZHEN TECHDOW	<u>1,000 UNITS/ML</u>	<u>A202732 001</u>	Jun 12, 2014
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HEPARIN SODIUM

	B BRAUN MEDICAL INC	5,000 UNITS/0.5ML	A208827 001	Nov 19, 2018
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+!	FRESENIUS KABI USA	10,000 UNITS/ML	N017029 020	Mar 31, 2011
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!	HOSPIRA	5,000 UNITS/ML	A088100 001	Apr 28, 1983
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+!	PFIZER	1,000 UNITS/ML	N201370 001	Jul 21, 2011
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+!		5,000 UNITS/ML	N201370 002	Jul 21, 2011
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+!		10,000 UNITS/ML	N201370 003	Jul 21, 2011
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HEPARIN SODIUM 12,500 UNITS IN DEXTROSE 5% IN PLASTIC CONTAINER

	HOSPIRA	5,000 UNITS/100ML	N019339 001	Mar 27, 1985
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HEPARIN SODIUM 12,500 UNITS IN SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER

	HOSPIRA	5,000 UNITS/100ML	N018916 006	Jan 31, 1984
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HEPARIN SODIUM 25,000 UNITS IN SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER

	HOSPIRA	5,000 UNITS/100ML	N018916 007	Jan 31, 1984
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		10,000 UNITS/100ML	N018916 008	Jan 31, 1984
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HEPARIN SODIUM PRESERVATIVE FREE

+!	FRESENIUS KABI USA	10,000 UNITS/ML	N017029 019	Nov 22, 2010
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!	HOSPIRA	10,000 UNITS/ML	A089522 001	May 04, 1987
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+!	PFIZER	1,000 UNITS/ML	N201370 004	Jul 21, 2011
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HEXACHLOROPHENE

SPONGE; TOPICAL

PRE-OP

<u>AT</u>	+! DAVIS AND GECK	<u>480MG</u>	<u>N017433 001</u>	
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PRE-OP II

<u>AT</u>	+ DAVIS AND GECK	<u>480MG</u>	<u>N017433 002</u>	
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HEXAMINOLEVULINATE HYDROCHLORIDE

FOR SOLUTION; INTRAVESICAL

CYSVIEW KIT

+!	PHOTOCURE ASA	100MG/VIAL	N022555 001	May 28, 2010
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HISTRELIN ACETATE

IMPLANT; SUBCUTANEOUS

SUPPRELIN LA

+!	ENDO PHARM	50MG	N022058 001	May 03, 2007
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VANTAS

+!	ENDO PHARM	50MG	N021732 001	Oct 12, 2004
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HOMATROPINE METHYLBROMIDE; HYDROCODONE BITARTRATE

SYRUP; ORAL

HYCODAN

<u>AA</u>	+ GENUS	<u>1.5MG/5ML; 5MG/5ML</u>	<u>N005213 002</u>	Jul 26, 1988
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HYDROCODONE BITARTRATE AND HOMATROPINE METHYLBROMIDE

<u>AA</u>	ABHAI LLC	<u>1.5MG/5ML; 5MG/5ML</u>	<u>A207487 001</u>	Feb 21, 2017
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<u>AA</u>	ACTAVIS MID	<u>1.5MG/5ML; 5MG/5ML</u>	<u>A088017 001</u>	Jul 05, 1983
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ATLANTIC

<u>AA</u>	! HI TECH PHARMA	<u>1.5MG/5ML; 5MG/5ML</u>	<u>A040613 001</u>	Feb 08, 2008
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<u>AA</u>	NOVEL LABS INC	<u>1.5MG/5ML; 5MG/5ML</u>	<u>A203535 001</u>	Feb 13, 2017
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<u>AA</u>	PADDOCK LLC	<u>1.5MG/5ML; 5MG/5ML</u>	<u>A205731 001</u>	Feb 15, 2017
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PRESCRIPTION DRUG PRODUCT LIST

HOMATROPINE METHYLBROMIDE; HYDROCODONE BITARTRATE

SYRUP; ORAL

HYDROCODONE BITARTRATE AND HOMATROPINE METHYLBROMIDE

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, 2017

AA ! NOVEL LABS INC **1.5MG; 5MG** **A091528 001** Apr 20, 2011

HYALURONIDASE

INJECTABLE; INJECTION

AMPHADASE

+! AMPHASTAR PHARM 150 UNITS/ML N021665 001 Oct 26, 2004

VITRASE

+! BAUSCH AND LOMB 200 UNITS/VIAL N021640 002 Dec 02, 2004

HYALURONIDASE RECOMBINANT HUMAN

INJECTABLE; INJECTION

HYLENEX RECOMBINANT

+! HALOZYME THERAP 150 UNITS/ML N021859 001 Dec 02, 2005

HYDRALAZINE HYDROCHLORIDE

INJECTABLE; INJECTION

HYDRALAZINE HYDROCHLORIDE

AP ! AKORN **20MG/ML** **A040730 001** Apr 21, 2009

AP FRESENIUS KABI USA **20MG/ML** **A040388 001** Mar 13, 2001

AP MYLAN INSTITUTIONAL **20MG/ML** **A204680 001** Apr 28, 2016

AP NAVINTA LLC **20MG/ML** **A202938 001** Mar 28, 2013

AP XGEN PHARMS **20MG/ML** **A203110 001** Jun 29, 2015

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 27, 2009

AA **25MG** **A090527 002** May 27, 2009

AA **50MG** **A090527 003** May 27, 2009

AA **100MG** **A090527 004** May 27, 2009

AA HERITAGE PHARMS INC **10MG** **A086242 001** Feb 04, 2010

AA **25MG** **A086242 003**

AA **50MG** **A086242 002**

AA **100MG** **A086242 004** Feb 04, 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 INVAGEN PHARMS **10MG** **A090255 001** Dec 15, 2008

AA **25MG** **A090255 002** Dec 15, 2008

AA **50MG** **A090255 003** Dec 15, 2008

AA **100MG** **A090255 004** Dec 15, 2008

AA PAR PHARM **10MG** **A087836 001** Oct 05, 1982

AA **25MG** **A086961 002**

AA **50MG** **A086962 001**

AA **100MG** **A088391 001** Sep 27, 1983

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

AA STRIDES PHARMA **25MG** **A200770 001** May 03, 2013

AA **50MG** **A200770 002** May 03, 2013

AA **100MG** **A200770 003** May 03, 2013

AA UPSHER SMITH LABS **10MG** **A209251 001** Jul 09, 2018

AA **25MG** **A209251 002** Jul 09, 2018

AA **50MG** **A209251 003** Jul 09, 2018

AA **100MG** **A209251 004** Jul 09, 2018

PRESCRIPTION DRUG PRODUCT LIST

HYDRALAZINE HYDROCHLORIDE; HYDROCHLOROTHIAZIDE

CAPSULE; ORAL

HYDRA-ZIDE

PAR PHARM	25MG; 25MG	A088957	001	Oct 21, 1985
!	50MG; 50MG	A088946	001	Oct 21, 1985

HYDRALAZINE HYDROCHLORIDE; ISOSORBIDE DINITRATE

TABLET; ORAL

BIDIL

+! ARBOR PHARMS LLC	37.5MG; 20MG	N020727	001	Jun 23, 2005
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HYDROCHLOROTHIAZIDE

CAPSULE; ORAL

HYDROCHLOROTHIAZIDE

AB	ALEMBIC PHARMS LTD	<u>12.5MG</u>	<u>A200645</u>	<u>001</u>	Nov 30, 2010
AB	AUROBINDO PHARMA	<u>12.5MG</u>	<u>A078164</u>	<u>001</u>	Sep 18, 2007
AB	IPCA LABS LTD	<u>12.5MG</u>	<u>A079237</u>	<u>001</u>	Apr 02, 2009
AB	IVAX SUB TEVA PHARMS	<u>12.5MG</u>	<u>A077005</u>	<u>001</u>	Jul 13, 2005
AB	JUBILANT CADISTA	<u>12.5MG</u>	<u>A078391</u>	<u>001</u>	Feb 11, 2008
AB	MYLAN	<u>12.5MG</u>	<u>A075640</u>	<u>001</u>	Jan 28, 2000
AB	PRINSTON INC	<u>12.5MG</u>	<u>A075907</u>	<u>001</u>	Sep 17, 2002
AB	SCIEGEN PHARMS INC	<u>12.5MG</u>	<u>A203561</u>	<u>001</u>	Jan 14, 2019
AB	SUN PHARM INDS INC	<u>12.5MG</u>	<u>A090651</u>	<u>001</u>	Apr 07, 2014
AB	UNICHEM	<u>12.5MG</u>	<u>A090510</u>	<u>001</u>	Jan 19, 2010

MICROZIDE

AB	+! ALLERGAN	<u>12.5MG</u>	<u>N020504</u>	<u>001</u>	Dec 27, 1996
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TABLET; ORAL

HYDROCHLOROTHIAZIDE

AB	!	ACCORD HLTHCARE	<u>12.5MG</u>	<u>A202556</u>	<u>001</u>	Sep 24, 2012
AB			<u>25MG</u>	<u>A202556</u>	<u>002</u>	Sep 24, 2012
AB			<u>50MG</u>	<u>A202556</u>	<u>003</u>	Sep 24, 2012
AB		ACTAVIS ELIZABETH	<u>12.5MG</u>	<u>A040707</u>	<u>001</u>	Feb 27, 2007
AB		AUROBINDO PHARMA	<u>25MG</u>	<u>A040780</u>	<u>001</u>	Jul 20, 2007
AB			<u>50MG</u>	<u>A040780</u>	<u>002</u>	Jul 20, 2007
AB		HERITAGE PHARMS INC	<u>25MG</u>	<u>A085182</u>	<u>002</u>	
AB			<u>50MG</u>	<u>A085182</u>	<u>001</u>	
AB		HIKMA INTL PHARMS	<u>50MG</u>	<u>A084878</u>	<u>001</u>	
AB		IPCA LABS LTD	<u>12.5MG</u>	<u>A040807</u>	<u>001</u>	Jul 20, 2007
AB			<u>25MG</u>	<u>A040807</u>	<u>002</u>	Jul 20, 2007
AB			<u>50MG</u>	<u>A040807</u>	<u>003</u>	Jul 20, 2007
AB		IVAX SUB TEVA PHARMS	<u>25MG</u>	<u>A083177</u>	<u>001</u>	
AB	!		<u>50MG</u>	<u>A083177</u>	<u>002</u>	
AB		LEADING PHARMA LLC	<u>12.5MG</u>	<u>A040702</u>	<u>003</u>	May 10, 2017
AB			<u>25MG</u>	<u>A040702</u>	<u>001</u>	Mar 16, 2007
AB			<u>50MG</u>	<u>A040702</u>	<u>002</u>	Mar 16, 2007
AB		OXFORD PHARMS	<u>25MG</u>	<u>A087059</u>	<u>001</u>	
AB			<u>50MG</u>	<u>A087068</u>	<u>001</u>	
AB		PRINSTON INC	<u>25MG</u>	<u>A040412</u>	<u>001</u>	Mar 29, 2002
AB			<u>50MG</u>	<u>A040412</u>	<u>002</u>	Mar 29, 2002
AB		SCIEGEN PHARMS INC	<u>25MG</u>	<u>A203018</u>	<u>001</u>	Jul 23, 2014
AB			<u>50MG</u>	<u>A203018</u>	<u>002</u>	Jul 23, 2014
AB		UNICHEM	<u>25MG</u>	<u>A040907</u>	<u>001</u>	Aug 15, 2008
AB			<u>50MG</u>	<u>A040907</u>	<u>002</u>	Aug 15, 2008

HYDROCHLOROTHIAZIDE; IRBESARTAN

TABLET; ORAL

AVALIDE

AB	+!	SANOFI AVENTIS US	<u>12.5MG; 150MG</u>	<u>N020758</u>	<u>002</u>	Sep 30, 1997
AB	+!		<u>12.5MG; 300MG</u>	<u>N020758</u>	<u>003</u>	Aug 31, 1998

IRBESARTAN AND HYDROCHLOROTHIAZIDE

AB		ALEMBIC PHARMS LTD	<u>12.5MG; 150MG</u>	<u>A091370</u>	<u>001</u>	Oct 15, 2012
AB			<u>12.5MG; 300MG</u>	<u>A091370</u>	<u>002</u>	Oct 15, 2012
AB			<u>25MG; 300MG</u>	<u>A091370</u>	<u>003</u>	Oct 12, 2016
AB		AUROBINDO PHARMA LTD	<u>12.5MG; 150MG</u>	<u>A203630</u>	<u>001</u>	Feb 22, 2013
AB			<u>12.5MG; 300MG</u>	<u>A203630</u>	<u>002</u>	Feb 22, 2013
AB			<u>25MG; 300MG</u>	<u>A203630</u>	<u>003</u>	Mar 31, 2016
AB		DR REDDYS LABS LTD	<u>12.5MG; 150MG</u>	<u>A203500</u>	<u>001</u>	Sep 27, 2012
AB			<u>12.5MG; 300MG</u>	<u>A203500</u>	<u>002</u>	Sep 27, 2012
AB		HIKMA	<u>12.5MG; 150MG</u>	<u>A090351</u>	<u>001</u>	Oct 15, 2012
AB			<u>12.5MG; 300MG</u>	<u>A090351</u>	<u>002</u>	Oct 15, 2012
AB			<u>25MG; 300MG</u>	<u>A090351</u>	<u>003</u>	Jun 08, 2017

PRESCRIPTION DRUG PRODUCT LISTHYDROCHLOROTHIAZIDE; IRBESARTAN

TABLET; ORAL

IRBESARTAN AND HYDROCHLOROTHIAZIDE

<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
<u>AB</u>	UNICHEM LABS LTD	<u>12.5MG;150MG</u>	<u>A207018 001</u>	Sep 19, 2017
<u>AB</u>		<u>12.5MG;300MG</u>	<u>A207018 002</u>	Sep 19, 2017

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>	HIKMA INTL PHARMS	<u>12.5MG;10MG</u>	<u>A076265 001</u>	Jul 08, 2002
<u>AB</u>		<u>12.5MG;20MG</u>	<u>A076265 002</u>	Jul 08, 2002
<u>AB</u>		<u>25MG;20MG</u>	<u>A076265 003</u>	Jul 08, 2002
<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>	+ ALVOGEN	<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>	+ MERCK SHARP DOHME	<u>12.5MG;100MG</u>	<u>N020387 003</u>	Oct 20, 2005
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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>	CADISTA PHARMS	<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>	HIKMA	<u>12.5MG;50MG</u>	<u>A077732 002</u>	Oct 06, 2010
<u>AB</u>		<u>12.5MG;100MG</u>	<u>A077732 001</u>	Apr 06, 2010
<u>AB</u>		<u>25MG;100MG</u>	<u>A077732 003</u>	Oct 06, 2010
<u>AB</u>	IPCA LABS LTD	<u>12.5MG;50MG</u>	<u>A201682 001</u>	Mar 01, 2013
<u>AB</u>		<u>12.5MG;100MG</u>	<u>A201682 002</u>	Mar 01, 2013
<u>AB</u>		<u>25MG;100MG</u>	<u>A201682 003</u>	Mar 01, 2013
<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

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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>	MYLAN	<u>12.5MG;50MG</u>	<u>A091652 001</u>	Oct 06, 2010
<u>AB</u>		<u>12.5MG;100MG</u>	<u>A091652 002</u>	Apr 06, 2010
<u>AB</u>		<u>25MG;100MG</u>	<u>A091652 003</u>	Oct 06, 2010
<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>	SANDOZ	<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>	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>	TORRENT PHARMS	<u>12.5MG;50MG</u>	<u>A090528 001</u>	Oct 06, 2010
<u>AB</u>		<u>12.5MG;100MG</u>	<u>A090528 003</u>	Apr 06, 2010
<u>AB</u>		<u>25MG;100MG</u>	<u>A090528 002</u>	Oct 06, 2010
<u>AB</u>	UNICHEM LABS LTD	<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; METHYLDOPA

TABLET; ORAL

METHYLDOPA AND HYDROCHLOROTHIAZIDE

	MYLAN	15MG;250MG	A070265 002	Jan 23, 1986
!		25MG;250MG	A070265 001	Jan 23, 1986

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

<u>AB</u>	+	US PHARMS HOLDINGS	<u>25MG;50MG</u>	<u>N018303 001</u>	Dec 31, 1984
		I			
<u>AB</u>	+	!	<u>25MG;100MG</u>	<u>N018303 002</u>	Dec 31, 1984
<u>AB</u>		ALEMBIC PHARMS LTD	<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
<u>AB</u>		SUN PHARM INDS	<u>25MG;50MG</u>	<u>A090654 001</u>	Jan 19, 2012
<u>AB</u>			<u>25MG;100MG</u>	<u>A090654 002</u>	Jan 19, 2012
<u>AB</u>			<u>50MG;100MG</u>	<u>A090654 003</u>	Jan 19, 2012

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>		HERITAGE PHARMS INC	<u>12.5MG;7.5MG</u>	<u>A202150 001</u>	Mar 07, 2014
<u>AB</u>			<u>12.5MG;15MG</u>	<u>A202150 002</u>	Mar 07, 2014
<u>AB</u>			<u>25MG;15MG</u>	<u>A202150 003</u>	Mar 07, 2014
<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

PRESCRIPTION DRUG PRODUCT LIST

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HYDROCHLOROTHIAZIDE; OLMESARTAN MEDOXOMIL

TABLET; ORAL

BENICAR HCT

<u>AB</u>	+	DAIICHI SANKYO	<u>12.5MG;20MG</u>	<u>N021532</u>	<u>002</u>	Jun 05, 2003
<u>AB</u>	+		<u>12.5MG;40MG</u>	<u>N021532</u>	<u>003</u>	Jun 05, 2003
<u>AB</u>	+	!	<u>25MG;40MG</u>	<u>N021532</u>	<u>005</u>	Jun 05, 2003

OLMESARTAN MEDOXOMIL AND HYDROCHLOROTHIAZIDE

<u>AB</u>		ALEMBIC PHARMS LTD	<u>12.5MG;20MG</u>	<u>A204233</u>	<u>001</u>	Apr 24, 2017
<u>AB</u>			<u>12.5MG;40MG</u>	<u>A204233</u>	<u>002</u>	Apr 24, 2017
<u>AB</u>			<u>25MG;40MG</u>	<u>A204233</u>	<u>003</u>	Apr 24, 2017
<u>AB</u>		AUROBINDO PHARMA LTD	<u>12.5MG;20MG</u>	<u>A205391</u>	<u>001</u>	Apr 24, 2017
<u>AB</u>			<u>12.5MG;40MG</u>	<u>A205391</u>	<u>002</u>	Apr 24, 2017
<u>AB</u>			<u>25MG;40MG</u>	<u>A205391</u>	<u>003</u>	Apr 24, 2017
<u>AB</u>		PRINSTON INC	<u>12.5MG;20MG</u>	<u>A207804</u>	<u>001</u>	Apr 24, 2017
<u>AB</u>			<u>12.5MG;40MG</u>	<u>A207804</u>	<u>002</u>	Apr 24, 2017
<u>AB</u>			<u>25MG;40MG</u>	<u>A207804</u>	<u>003</u>	Apr 24, 2017
<u>AB</u>		TEVA PHARMS USA	<u>12.5MG;40MG</u>	<u>A200532</u>	<u>002</u>	Apr 24, 2017
<u>AB</u>		TORRENT	<u>12.5MG;20MG</u>	<u>A206515</u>	<u>001</u>	May 03, 2017
<u>AB</u>			<u>12.5MG;40MG</u>	<u>A206515</u>	<u>002</u>	May 03, 2017
<u>AB</u>			<u>25MG;40MG</u>	<u>A206515</u>	<u>003</u>	May 03, 2017
<u>AB</u>		UMEDICA LABS PVT LTD	<u>12.5MG;20MG</u>	<u>A208847</u>	<u>001</u>	Sep 17, 2019
<u>AB</u>			<u>12.5MG;40MG</u>	<u>A208847</u>	<u>003</u>	Sep 17, 2019
<u>AB</u>			<u>25MG;40MG</u>	<u>A208847</u>	<u>002</u>	Sep 17, 2019

HYDROCHLOROTHIAZIDE; PROPRANOLOL HYDROCHLORIDE

TABLET; ORAL

PROPRANOLOL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE

<u>AB</u>	!	MYLAN	<u>25MG;80MG</u>	<u>A070947</u>	<u>001</u>	Apr 01, 1987
	!		<u>25MG;40MG</u>	<u>A070947</u>	<u>002</u>	Mar 04, 1987

HYDROCHLOROTHIAZIDE; QUINAPRIL HYDROCHLORIDE

TABLET; ORAL

ACCURETIC

<u>AB</u>	+	PFIZER PHARMS	<u>12.5MG;EQ 10MG BASE</u>	<u>N020125</u>	<u>001</u>	Dec 28, 1999
<u>AB</u>	+		<u>12.5MG;EQ 20MG BASE</u>	<u>N020125</u>	<u>002</u>	Dec 28, 1999
<u>AB</u>	+	!	<u>25MG;EQ 20MG BASE</u>	<u>N020125</u>	<u>003</u>	Dec 28, 1999

QUINAPRIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE

<u>AB</u>		APOTEX CORP	<u>12.5MG;EQ 10MG BASE</u>	<u>A091524</u>	<u>001</u>	Mar 12, 2013
<u>AB</u>			<u>12.5MG;EQ 20MG BASE</u>	<u>A091524</u>	<u>002</u>	Mar 12, 2013
<u>AB</u>			<u>25MG;EQ 20MG BASE</u>	<u>A091524</u>	<u>003</u>	Mar 12, 2013
<u>AB</u>		AUROBINDO PHARMA	<u>12.5MG;EQ 10MG BASE</u>	<u>A078450</u>	<u>001</u>	Aug 24, 2007
<u>AB</u>			<u>12.5MG;EQ 20MG BASE</u>	<u>A078450</u>	<u>002</u>	Aug 24, 2007
<u>AB</u>			<u>25MG;EQ 20MG BASE</u>	<u>A078450</u>	<u>003</u>	Aug 24, 2007
<u>AB</u>		INVAGEN PHARMS	<u>12.5MG;EQ 10MG BASE</u>	<u>A201356</u>	<u>001</u>	Apr 20, 2011
<u>AB</u>			<u>12.5MG;EQ 20MG BASE</u>	<u>A201356</u>	<u>002</u>	Apr 20, 2011
<u>AB</u>			<u>25MG;EQ 20MG BASE</u>	<u>A201356</u>	<u>003</u>	Apr 20, 2011
<u>AB</u>		LUPIN	<u>12.5MG;EQ 10MG BASE</u>	<u>A076374</u>	<u>001</u>	Mar 31, 2004
<u>AB</u>			<u>12.5MG;EQ 20MG BASE</u>	<u>A076374</u>	<u>002</u>	Mar 31, 2004
<u>AB</u>			<u>25MG;EQ 20MG BASE</u>	<u>A076374</u>	<u>003</u>	Mar 31, 2004

HYDROCHLOROTHIAZIDE; SPIRONOLACTONE

TABLET; ORAL

ALDACTAZIDE

<u>AB</u>	+	GD SEARLE LLC	<u>25MG;25MG</u>	<u>N012616</u>	<u>004</u>	Dec 30, 1982
<u>AB</u>		MYLAN	<u>25MG;25MG</u>	<u>A086513</u>	<u>001</u>	
<u>AB</u>		SUN PHARM INDUSTRIES	<u>25MG;25MG</u>	<u>A089534</u>	<u>001</u>	Jul 02, 1987
		ALDACTAZIDE				
	+	GD SEARLE LLC	<u>50MG;50MG</u>	<u>N012616</u>	<u>005</u>	Dec 30, 1982

HYDROCHLOROTHIAZIDE; TELMISARTAN

TABLET; ORAL

MICARDIS HCT

<u>AB</u>	+	BOEHRINGER INGELHEIM	<u>12.5MG;40MG</u>	<u>N021162</u>	<u>001</u>	Nov 17, 2000
<u>AB</u>	+		<u>12.5MG;80MG</u>	<u>N021162</u>	<u>002</u>	Nov 17, 2000
<u>AB</u>	+	!	<u>25MG;80MG</u>	<u>N021162</u>	<u>003</u>	Apr 19, 2004

TELMISARTAN AND HYDROCHLOROTHIAZIDE

<u>AB</u>		ALEMBIC PHARMS LTD	<u>12.5MG;40MG</u>	<u>A203010</u>	<u>001</u>	Feb 25, 2014
<u>AB</u>			<u>12.5MG;80MG</u>	<u>A203010</u>	<u>002</u>	Feb 25, 2014
<u>AB</u>			<u>25MG;80MG</u>	<u>A203010</u>	<u>003</u>	Feb 25, 2014

PRESCRIPTION DRUG PRODUCT LIST

3-225 (of 453)

HYDROCHLOROTHIAZIDE; TELMISARTAN

TABLET; ORAL

TELMISARTAN AND HYDROCHLOROTHIAZIDE

AB	AUROBINDO PHARMA LTD	12.5MG;40MG	A208727 001	Dec 15, 2016
AB		12.5MG;80MG	A208727 002	Dec 15, 2016
AB		25MG;80MG	A208727 003	Dec 15, 2016
AB	GLENMARK PHARMS LTD	12.5MG;40MG	A202544 001	Mar 04, 2019
AB		12.5MG;80MG	A202544 002	Mar 04, 2019
AB		25MG;80MG	A202544 003	Mar 04, 2019
AB	LUPIN LTD	12.5MG;40MG	A091351 001	Aug 07, 2014
AB		12.5MG;80MG	A091351 002	Aug 07, 2014
AB		25MG;80MG	A091351 003	Aug 07, 2014
AB	MACLEODS PHARMS LTD	12.5MG;40MG	A204169 001	Nov 02, 2015
AB		12.5MG;80MG	A204169 002	Nov 02, 2015
AB		25MG;80MG	A204169 003	Nov 02, 2015
AB	MYLAN	12.5MG;40MG	A091648 001	Feb 25, 2014
AB		12.5MG;80MG	A091648 002	Feb 25, 2014
AB		25MG;80MG	A091648 003	Feb 25, 2014
AB	PRINSTON INC	12.5MG;40MG	A209028 001	Nov 06, 2017
AB		12.5MG;80MG	A209028 002	Nov 06, 2017
AB		25MG;80MG	A209028 003	Nov 06, 2017
AB	TORRENT	12.5MG;40MG	A201192 001	Feb 25, 2014
AB		12.5MG;80MG	A201192 002	Feb 25, 2014
AB		25MG;80MG	A201192 003	Feb 25, 2014
AB	ZYDUS PHARMS	12.5MG;40MG	A204221 001	Aug 15, 2017
AB		12.5MG;80MG	A204221 002	Aug 15, 2017
AB		25MG;80MG	A204221 003	Aug 15, 2017

HYDROCHLOROTHIAZIDE; TRIAMTERENE

CAPSULE; ORAL

DYAZIDE

AB	+ GLAXOSMITHKLINE LLC	25MG;37.5MG	N016042 003	Mar 03, 1994
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TRIAMTERENE AND HYDROCHLOROTHIAZIDE

AB	CADILA	25MG;37.5MG	A208358 001	Feb 11, 2019
AB	DURAMED PHARMS BARR	25MG;37.5MG	A075052 001	Jun 18, 1999
AB	LANNETT CO INC	25MG;37.5MG	A201407 001	Dec 09, 2011
AB	MYLAN	25MG;37.5MG	A074701 001	Jun 07, 1996
AB	SANDOZ	25MG;37.5MG	A074821 001	Jun 05, 1997

TABLET; ORAL

MAXZIDE

AB	+ MYLAN PHARMS INC	50MG;75MG	N019129 001	Oct 22, 1984
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MAXZIDE-25

AB	+ MYLAN PHARMS INC	25MG;37.5MG	N019129 003	May 13, 1988
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TRIAMTERENE AND HYDROCHLOROTHIAZIDE

AB	APOTEX INC	25MG;37.5MG	A071251 002	May 05, 1998
AB		50MG;75MG	A071251 001	Apr 17, 1988
AB	PLIVA	25MG;37.5MG	A074026 001	Apr 26, 1996
AB	SANDOZ	25MG;37.5MG	A073281 001	Apr 30, 1992
AB		50MG;75MG	A072011 001	Jun 17, 1988
AB	WATSON LABS	25MG;37.5MG	A073449 001	Sep 23, 1993
AB		50MG;75MG	A071851 001	Nov 30, 1988
AB	ZYDUS PHARMS	25MG;37.5MG	A208360 001	Jun 29, 2018
AB		50MG;75MG	A208360 002	Jun 29, 2018

HYDROCHLOROTHIAZIDE; VALSARTAN

TABLET; ORAL

DIOVAN HCT

AB	+ NOVARTIS	12.5MG;80MG	N020818 001	Mar 06, 1998
AB	+	12.5MG;160MG	N020818 002	Mar 06, 1998
AB	+	12.5MG;320MG	N020818 004	Apr 28, 2006
AB	+	25MG;160MG	N020818 003	Jan 17, 2002
AB	+ !	25MG;320MG	N020818 005	Apr 28, 2006

VALSARTAN AND HYDROCHLOROTHIAZIDE

AB	ALEMBIC PHARMS LTD	12.5MG;80MG	A201662 001	Mar 21, 2013
AB		12.5MG;160MG	A201662 002	Mar 21, 2013
AB		12.5MG;320MG	A201662 003	Mar 21, 2013
AB		25MG;160MG	A201662 004	Mar 21, 2013
AB		25MG;320MG	A201662 005	Mar 21, 2013
AB	AUROBINDO PHARMA LTD	12.5MG;80MG	A202519 001	Mar 21, 2013
AB		12.5MG;160MG	A202519 002	Mar 21, 2013
AB		12.5MG;320MG	A202519 003	Mar 21, 2013
AB		25MG;160MG	A202519 004	Mar 21, 2013

PRESCRIPTION DRUG PRODUCT LIST

HYDROCHLOROTHIAZIDE; VALSARTAN

TABLET; ORAL

VALSARTAN AND HYDROCHLOROTHIAZIDE

<u>AB</u>		<u>25MG; 320MG</u>	<u>A202519 005</u>	Mar 21, 2013
<u>AB</u>	LUPIN LTD	<u>12.5MG; 80MG</u>	<u>A078946 003</u>	Mar 21, 2013
<u>AB</u>		<u>12.5MG; 160MG</u>	<u>A078946 004</u>	Mar 21, 2013
<u>AB</u>		<u>12.5MG; 320MG</u>	<u>A078946 001</u>	Mar 21, 2013
<u>AB</u>		<u>25MG; 160MG</u>	<u>A078946 005</u>	Mar 21, 2013
<u>AB</u>		<u>25MG; 320MG</u>	<u>A078946 002</u>	Mar 21, 2013
<u>AB</u>	MACLEODS PHARMS LTD	<u>12.5MG; 80MG</u>	<u>A203145 001</u>	Apr 19, 2013
<u>AB</u>		<u>12.5MG; 160MG</u>	<u>A203145 002</u>	Apr 19, 2013
<u>AB</u>		<u>12.5MG; 320MG</u>	<u>A203145 003</u>	Apr 19, 2013
<u>AB</u>		<u>25MG; 160MG</u>	<u>A203145 004</u>	Apr 19, 2013
<u>AB</u>		<u>25MG; 320MG</u>	<u>A203145 005</u>	Apr 19, 2013
<u>AB</u>	MYLAN PHARMS INC	<u>12.5MG; 80MG</u>	<u>A078020 001</u>	Sep 21, 2012
<u>AB</u>		<u>12.5MG; 160MG</u>	<u>A078020 002</u>	Sep 21, 2012
<u>AB</u>		<u>12.5MG; 320MG</u>	<u>A078020 004</u>	Sep 21, 2012
<u>AB</u>		<u>25MG; 160MG</u>	<u>A078020 003</u>	Sep 21, 2012
<u>AB</u>		<u>25MG; 320MG</u>	<u>A078020 005</u>	Sep 21, 2012
<u>AB</u>	PRINSTON INC	<u>12.5MG; 80MG</u>	<u>A206083 001</u>	Feb 08, 2016
<u>AB</u>		<u>12.5MG; 160MG</u>	<u>A206083 002</u>	Feb 08, 2016
<u>AB</u>		<u>12.5MG; 320MG</u>	<u>A206083 003</u>	Feb 08, 2016
<u>AB</u>		<u>25MG; 160MG</u>	<u>A206083 004</u>	Feb 08, 2016
<u>AB</u>		<u>25MG; 320MG</u>	<u>A206083 005</u>	Feb 08, 2016

HYDROCODONE BITARTRATE

CAPSULE, EXTENDED RELEASE; ORAL

HYDROCODONE BITARTRATE

<u>AB</u>	ALVOGEN	<u>10MG</u>	<u>A206986 001</u>	Jan 21, 2020
<u>AB</u>		<u>15MG</u>	<u>A206986 002</u>	Jan 21, 2020
<u>AB</u>		<u>20MG</u>	<u>A206986 003</u>	Jan 21, 2020
<u>AB</u>		<u>30MG</u>	<u>A206986 004</u>	Jan 21, 2020
<u>AB</u>		<u>40MG</u>	<u>A206986 005</u>	Jan 21, 2020
<u>AB</u>		<u>50MG</u>	<u>A206986 006</u>	Jan 21, 2020
	<u>ZOHYDRO ER</u>			
<u>AB</u>	+! PERSION	<u>10MG</u>	<u>N202880 001</u>	Oct 25, 2013
<u>AB</u>	+	<u>15MG</u>	<u>N202880 002</u>	Oct 25, 2013
<u>AB</u>	+	<u>20MG</u>	<u>N202880 003</u>	Oct 25, 2013
<u>AB</u>	+	<u>30MG</u>	<u>N202880 004</u>	Oct 25, 2013
<u>AB</u>	+	<u>40MG</u>	<u>N202880 005</u>	Oct 25, 2013
<u>AB</u>	+	<u>50MG</u>	<u>N202880 006</u>	Oct 25, 2013

TABLET, EXTENDED RELEASE; ORAL

HYSINGLA ER

+!	PURDUE PHARMA LP	20MG	N206627 001	Nov 20, 2014
+		30MG	N206627 002	Nov 20, 2014
+		40MG	N206627 003	Nov 20, 2014
+		60MG	N206627 004	Nov 20, 2014
+		80MG	N206627 005	Nov 20, 2014
+		100MG	N206627 006	Nov 20, 2014
+		120MG	N206627 007	Nov 20, 2014

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>5MG; 200MG</u>	<u>A076642 002</u>	Mar 18, 2004
<u>AB</u>	!	<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
<u>AB</u>	VINTAGE PHARMS	<u>5MG; 200MG</u>	<u>A077727 001</u>	Nov 06, 2006
<u>AB</u>		<u>7.5MG; 200MG</u>	<u>A077723 001</u>	Nov 06, 2006
<u>AB</u>		<u>10MG; 200MG</u>	<u>A077723 002</u>	Nov 06, 2006
	<u>REPREXAIN</u>			
<u>AB</u>	AMNEAL PHARMS NY	<u>10MG; 200MG</u>	<u>A076642 004</u>	Oct 19, 2007
		<u>2.5MG; 200MG</u>	<u>A076642 003</u>	Oct 19, 2007

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>	
	<u>ANUSOL HC</u>			
<u>AT</u>	SALIX PHARMS	<u>2.5%</u>	<u>A088250 001</u>	Jun 06, 1984

PRESCRIPTION DRUG PRODUCT LIST

HYDROCORTISONE

CREAM; TOPICAL

HYDROCORTISONE

AT	ACTAVIS MID ATLANTIC	1%	A087795 001	May 03, 1983
AT		2.5%	A089682 001	Mar 10, 1988
AT	! FOUGERA PHARMS INC	1%	A080693 003	
AT	! FOUGERA PHARMS INC	2.5%	A089414 001	Dec 16, 1986
AT	LANNETT CO INC	2.5%	A040503 001	Mar 12, 2004
AT	PERRIGO NEW YORK	2.5%	A085025 001	
AT	RISING	2.5%	A040879 001	Aug 20, 2010
AT	TARO PHARM INDS LTD	2.5%	A088799 001	Nov 09, 1984
AT	TELOGENT PHARMA INC	2.5%	A203810 001	Jul 23, 2018

ENEMA; RECTAL

COLOCORT

AB	PADDOCK LLC	100MG/60ML	A075172 001	Dec 03, 1999
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CORTENEMA

AB	+! ANI PHARMS	100MG/60ML	N016199 001	
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HYDROCORTISONE

AB	TEVA PHARMS	100MG/60ML	A074171 001	May 27, 1994
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LOTION; TOPICAL

HYDROCORTISONE

AT	! FOUGERA PHARMS	2.5%	A040351 001	Jul 25, 2000
AT	LANNETT CO INC	2.5%	A040417 001	Jul 30, 2003
AT	TARO	2.5%	A040247 001	Jul 23, 1999
AT	TELOGENT PHARMA INC	2.5%	A203804 001	Jul 27, 2018

STIE-CORT

AT	PERRIGO CO	2.5%	A089074 001	Nov 26, 1985
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ALA-SCALP

	MARNEL PHARMS	2%	A083231 001	
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OINTMENT; TOPICAL

HYDROCORTISONE

AT	ACTAVIS MID ATLANTIC	1%	A087796 001	Oct 13, 1982
AT	! FOUGERA PHARMS	1%	A080692 001	
AT	! FOUGERA PHARMS INC	2.5%	A081203 001	May 28, 1993
AT	PERRIGO NEW YORK	2.5%	A085027 001	
AT	TARO	1%	A086257 001	

HYDROCORTISONE IN ABSORBASE

AT	CMP PHARMA INC	1%	A088138 001	Sep 06, 1985
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SOLUTION; TOPICAL

TEXACORT

!	MISSION PHARMA	2.5%	A081271 001	Apr 17, 1992
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TABLET; ORAL

CORTEF

AB	+ PHARMACIA AND UPJOHN	5MG	N008697 003	
AB	+	10MG	N008697 001	
AB	+!	20MG	N008697 002	

HYDROCORTISONE

AB	HIKMA INTL PHARMS	5MG	A083365 002	Feb 23, 2015
AB		10MG	A083365 003	Feb 23, 2015
AB		20MG	A083365 001	
AB	IMPAX LABS INC	5MG	A040646 001	Mar 30, 2007
AB		10MG	A040646 002	Mar 30, 2007
AB		20MG	A040646 003	Mar 30, 2007
AB	STRIDES PHARMA	5MG	A207029 001	Apr 27, 2017
AB		10MG	A207029 002	Apr 27, 2017
AB		20MG	A207029 003	Apr 27, 2017
AB	VINTAGE	5MG	A040761 001	Jul 16, 2007
AB		10MG	A040761 002	Jul 16, 2007
AB		20MG	A040761 003	Jul 16, 2007

HYDROCORTISONE ACETATE

AEROSOL, METERED; RECTAL

CORTIFOAM

+	MYLAN SPECIALITY LP	10%	N017351 001	Feb 10, 1982
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CREAM; TOPICAL

MICORT-HC

	SEBELA IRELAND LTD	2.5%	A040396 001	Feb 27, 2001
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PRESCRIPTION DRUG PRODUCT LIST

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; 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

SEBELA IRELAND LTD 0.5%;1% A083778 001

1%;1% A085368 001

LOTION; TOPICAL

PRAMOSONE

SEBELA IRELAND LTD 1%;1% A085980 001

2.5%;1% A085979 001

HYDROCORTISONE ACETATE; UREA

CREAM; TOPICAL

U-CORT

TARO 1%;10% A089472 001 Jun 13, 1988

HYDROCORTISONE BUTYRATE

CREAM; TOPICAL

HYDROCORTISONE BUTYRATE

AB1 TARO PHARM INDS **0.1%** **A076654 001** Aug 03, 2005

LOCOID

AB1 +! BAUSCH **0.1%** **N018514 001** Mar 31, 1982

HYDROCORTISONE BUTYRATE

AB2 ACTAVIS MID **0.1%** **A205134 001** Dec 08, 2017

ATLANTIC

AB2 GLENMARK GENERICS **0.1%** **A202145 001** Sep 27, 2013

LOCOID 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 TELIGENT PHARMA INC **0.1%** **A209556 001** Nov 21, 2017

LOCOID

AB +! BAUSCH **0.1%** **N022076 001** May 18, 2007

OINTMENT; TOPICAL

HYDROCORTISONE BUTYRATE

AB TARO **0.1%** **A076842 001** Dec 27, 2004

LOCOID

AB +! PRECISION DERMAT **0.1%** **N018652 001** Oct 29, 1982

SOLUTION; TOPICAL

HYDROCORTISONE BUTYRATE

AT TARO PHARM INDS **0.1%** **A076364 001** Jan 14, 2004

LOCOID

AT +! BAUSCH **0.1%** **N019116 001** Feb 25, 1987

HYDROCORTISONE PROBUTATE

CREAM; TOPICAL

PANDEL

+! FOUGERA PHARMS 0.1% N020453 001 Feb 28, 1997

HYDROCORTISONE SODIUM SUCCINATE

INJECTABLE; INJECTION

SOLU-CORTEF

+! 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 GLENMARK PHARMS LTD **0.2%** **A211129 001** Oct 12, 2018

AB LUPIN LTD **0.2%** **A210307 001** Aug 15, 2019

AB PERRIGO ISRAEL **0.2%** **A075666 001** May 24, 2000

AB ! TARO **0.2%** **A075042 001** Aug 25, 1998

PRESCRIPTION DRUG PRODUCT LIST

HYDROCORTISONE VALERATE

OINTMENT; TOPICAL

HYDROCORTISONE VALERATE

<u>AB</u>	COSETTE	<u>0.2%</u>	<u>A211764</u>	<u>001</u>	Mar 04, 2020
<u>AB</u>	GLENMARK PHARMS LTD	<u>0.2%</u>	<u>A211750</u>	<u>001</u>	Dec 14, 2018
<u>AB</u>	! TARO	<u>0.2%</u>	<u>A075043</u>	<u>001</u>	Aug 25, 1998

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</u>	<u>001</u>	Dec 29, 1995
<u>AT</u>	SANDOZ INC	<u>1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML</u>	<u>A062423</u>	<u>001</u>	Aug 25, 1983

SUSPENSION/DROPS; OPHTHALMIC

NEOMYCIN AND POLYMYXIN B SULFATES AND HYDROCORTISONE

!	SANDOZ INC	1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML	A062874	001	May 11, 1988
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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</u>	<u>001</u>	
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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</u>	<u>001</u>	May 01, 2006
<u>AT</u>	SANDOZ INC	<u>1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML</u>	<u>A062488</u>	<u>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</u>	<u>001</u>	Aug 28, 1996
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HYDROMORPHONE HYDROCHLORIDE

INJECTABLE; INJECTION

DILAUDID

<u>AP</u>	+! FRESENIUS KABI USA	<u>1MG/ML</u>	<u>N019034</u>	<u>003</u>	Apr 30, 2009
<u>AP</u>	+!	<u>2MG/ML</u>	<u>N019034</u>	<u>004</u>	Apr 30, 2009

HYDROMORPHONE HYDROCHLORIDE

<u>AP</u>	AKORN	<u>10MG/ML</u>	<u>A078228</u>	<u>001</u>	Apr 14, 2010
<u>AP</u>		<u>10MG/ML</u>	<u>A078261</u>	<u>001</u>	Apr 14, 2010
<u>AP</u>	BARR	<u>10MG/ML</u>	<u>A076444</u>	<u>001</u>	Apr 25, 2003
<u>AP</u>	EUROHLTH INTL SARL	<u>2MG/ML</u>	<u>A202159</u>	<u>001</u>	Apr 27, 2018
<u>AP</u>	HOSPIRA INC	<u>1MG/ML</u>	<u>N200403</u>	<u>001</u>	Dec 01, 2011
<u>AP</u>		<u>2MG/ML</u>	<u>N200403</u>	<u>002</u>	Dec 01, 2011
<u>AP</u>		<u>4MG/ML</u>	<u>N200403</u>	<u>003</u>	Dec 01, 2011
<u>AP</u>		<u>10MG/ML</u>	<u>A078591</u>	<u>001</u>	Jun 17, 2008

DILAUDID

+!	FRESENIUS KABI USA	0.2MG/ML	N019034	006	Jan 16, 2020
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SOLUTION; ORAL

DILAUDID

<u>AA</u>	+! RHODES PHARMS	<u>5MG/5ML</u>	<u>N019891</u>	<u>001</u>	Dec 07, 1992
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HYDROMORPHONE HYDROCHLORIDE

<u>AA</u>	ASCENT PHARMS INC	<u>5MG/5ML</u>	<u>A210176</u>	<u>001</u>	Oct 27, 2017
<u>AA</u>	HIKMA	<u>5MG/5ML</u>	<u>A074653</u>	<u>001</u>	Jul 29, 1998

TABLET; ORAL

DILAUDID

<u>AB</u>	+ RHODES PHARMS	<u>2MG</u>	<u>N019892</u>	<u>003</u>	Nov 09, 2007
<u>AB</u>	+	<u>4MG</u>	<u>N019892</u>	<u>002</u>	Nov 09, 2007
<u>AB</u>	+!	<u>8MG</u>	<u>N019892</u>	<u>001</u>	Dec 07, 1992

HYDROMORPHONE HYDROCHLORIDE

<u>AB</u>	ASCENT PHARMS INC	<u>2MG</u>	<u>A210506</u>	<u>001</u>	Jan 17, 2018
<u>AB</u>		<u>4MG</u>	<u>A210506</u>	<u>002</u>	Jan 17, 2018
<u>AB</u>		<u>8MG</u>	<u>A210506</u>	<u>003</u>	Jan 17, 2018
<u>AB</u>	AUROLIFE PHARMA LLC	<u>2MG</u>	<u>A205814</u>	<u>001</u>	May 13, 2016
<u>AB</u>		<u>4MG</u>	<u>A205814</u>	<u>002</u>	May 13, 2016
<u>AB</u>		<u>8MG</u>	<u>A205814</u>	<u>003</u>	May 13, 2016
<u>AB</u>	HIKMA	<u>4MG</u>	<u>A074597</u>	<u>003</u>	May 29, 2009
<u>AB</u>		<u>8MG</u>	<u>A074597</u>	<u>001</u>	Jul 29, 1998
<u>AB</u>	LANNETT CO INC	<u>2MG</u>	<u>A077471</u>	<u>002</u>	Dec 09, 2009
<u>AB</u>		<u>4MG</u>	<u>A077471</u>	<u>003</u>	Dec 09, 2009
<u>AB</u>		<u>8MG</u>	<u>A077471</u>	<u>001</u>	Dec 09, 2009
<u>AB</u>	NOSTRUM LABS INC	<u>8MG</u>	<u>A076723</u>	<u>001</u>	Oct 18, 2005
<u>AB</u>	SPECGX LLC	<u>2MG</u>	<u>A076855</u>	<u>002</u>	Sep 19, 2007
<u>AB</u>		<u>4MG</u>	<u>A076855</u>	<u>003</u>	Sep 19, 2007
<u>AB</u>		<u>8MG</u>	<u>A076855</u>	<u>001</u>	Dec 23, 2004

TABLET, EXTENDED RELEASE; ORAL

HYDROMORPHONE HYDROCHLORIDE

<u>AB</u>	OSMOTICA	<u>8MG</u>	<u>A205629</u>	<u>001</u>	Jul 07, 2016
<u>AB</u>		<u>12MG</u>	<u>A205629</u>	<u>002</u>	Jul 07, 2016
<u>AB</u>		<u>16MG</u>	<u>A205629</u>	<u>003</u>	Jul 07, 2016
<u>AB</u>	!	<u>32MG</u>	<u>A205629</u>	<u>004</u>	Jul 07, 2016

PRESCRIPTION DRUG PRODUCT LIST

HYDROMORPHONE HYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL

HYDROMORPHONE HYDROCHLORIDE

AB	PADDOCK LLC	8MG	A204278 001	Apr 06, 2015
AB		12MG	A204278 002	Apr 06, 2015
AB		16MG	A204278 003	Apr 06, 2015
AB		32MG	A204278 004	Sep 20, 2017

HYDROXOCOBALAMIN

INJECTABLE; INJECTION

CYANOKIT

+! SERB SA 5GM/VIAL (5GM/KIT) N022041 001 Apr 08, 2011

HYDROXOCOBALAMIN

! ACTAVIS LLC 1MG/ML A085998 001

HYDROXYAMPHETAMINE HYDROBROMIDE; TROPICAMIDE

SOLUTION/DROPS;OPHTHALMIC

PAREMYD

+! AKORN 1%;0.25% N019261 001 Jan 30, 1992

HYDROXYCHLOROQUINE SULFATE

TABLET;ORAL

HYDROXYCHLOROQUINE SULFATE

AB	ALKALOIDA ZRT	200MG	A201691 001	May 08, 2018
AB	AMNEAL PHARMS CO	200MG	A210577 001	May 15, 2018
AB	HIKMA PHARMS	200MG	A040760 001	Aug 15, 2007
AB	IPCA LABS LTD	200MG	A040766 001	Jun 14, 2007
AB	MYLAN	200MG	A040274 001	May 29, 1998
AB	SANDOZ	200MG	A040104 001	Nov 30, 1995
AB	TEVA PHARMS	200MG	A040081 001	Sep 30, 1994
AB	TWI PHARMS	200MG	A210441 001	May 01, 2018
AB	WATSON LABS	200MG	A040133 001	Nov 30, 1995
AB	ZYDUS PHARMS USA INC	200MG	A040657 001	Sep 21, 2007

PLAQUENIL**AB** +! CONCORDIA **200MG** **N009768 001**HYDROXYPROGESTERONE CAPROATE

SOLUTION;INTRAMUSCULAR

HYDROXYPROGESTERONE CAPROATE

AP	AM REGENT	250MG/ML (250MG/ML)	A210723 001	Jun 21, 2018
AP	ASPEN	250MG/ML (250MG/ML)	A211777 001	Aug 08, 2019
AP	EUGIA PHARMA	250MG/ML (250MG/ML)	A211071 001	Apr 16, 2019
AP	SLAYBACK PHARMA LLC	250MG/ML (250MG/ML)	A210877 001	Mar 22, 2019

MAKENA PRESERVATIVE FREE**AP** +! AMAG PHARMA USA **250MG/ML (250MG/ML)** **N021945 002** Feb 19, 2016HYDROXYPROGESTERONE CAPROATE

AP1	EUGIA PHARMA	1250MG/5ML (250MG/ML)	A211070 001	Apr 16, 2019
AP1	SLAYBACK PHARMA LLC	1250MG/5ML (250MG/ML)	A210618 001	Dec 28, 2018
AP1	SUN PHARM	1250MG/5ML (250MG/ML)	A208381 001	Apr 09, 2019

MAKENA**AP1** +! AMAG PHARMA USA **1250MG/5ML (250MG/ML)** **N021945 001** Feb 03, 2011

HYDROXYPROGESTERONE CAPROATE

! ASPEN GLOBAL INC 1250MG/5ML (250MG/ML) A200271 001 Aug 24, 2015

SOLUTION;SUBCUTANEOUS

MAKENA (AUTOINJECTOR)

+! AMAG PHARMA USA 275MG/1.1ML (250MG/ML) N021945 004 Feb 14, 2018

HYDROXYPROPYL CELLULOSE

INSERT;OPHTHALMIC

LACRISERT

+! VALEANT PHARMS INTL 5MG N018771 001

HYDROXYUREA

CAPSULE;ORAL

HYDREA**AB** +! BRISTOL MYERS
SQUIBB **500MG** **N016295 001**HYDROXYUREA**AB** BARR **500MG** **A075143 001** Oct 16, 1998**AB** PAR PHARM **500MG** **A075340 001** Feb 24, 1999

DROXIA

+ BRISTOL MYERS 200MG N016295 002 Feb 25, 1998

SQUIBB

+ 300MG N016295 003 Feb 25, 1998

+ 400MG N016295 004 Feb 25, 1998

PRESCRIPTION DRUG PRODUCT LIST

HYDROXYUREA

TABLET; ORAL

SIKLOS

+	ADDMEDICA SAS	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

AA	!	HI TECH PHARMA	<u>10MG/5ML</u>	<u>A040010</u>	<u>001</u>	Oct 28, 1994
AA		LANNETT CO INC	<u>10MG/5ML</u>	<u>A201674</u>	<u>001</u>	Aug 21, 2013
AA	!	VINTAGE PHARMS	<u>10MG/5ML</u>	<u>A040391</u>	<u>001</u>	Apr 10, 2002
AA	!	WOCKHARDT BIO AG	<u>10MG/5ML</u>	<u>A087294</u>	<u>001</u>	Apr 12, 1982

TABLET; ORAL

HYDROXYZINE HYDROCHLORIDE

AB		AMNEAL PHARM	<u>10MG</u>	<u>A040808</u>	<u>001</u>	Sep 24, 2008
AB			<u>25MG</u>	<u>A040808</u>	<u>002</u>	Sep 24, 2008
AB			<u>50MG</u>	<u>A040808</u>	<u>003</u>	Sep 24, 2008
AB		ECI PHARMS LLC	<u>10MG</u>	<u>A040804</u>	<u>001</u>	Jun 30, 2008
AB			<u>25MG</u>	<u>A040804</u>	<u>002</u>	Jun 30, 2008
AB			<u>50MG</u>	<u>A040804</u>	<u>003</u>	Jun 30, 2008
AB		EPIC PHARMA LLC	<u>10MG</u>	<u>A040604</u>	<u>002</u>	Dec 28, 2004
AB			<u>25MG</u>	<u>A040604</u>	<u>003</u>	Dec 28, 2004
AB			<u>50MG</u>	<u>A040604</u>	<u>001</u>	Dec 28, 2004
AB		HERITAGE PHARMA	<u>10MG</u>	<u>A204279</u>	<u>001</u>	Aug 20, 2014
AB			<u>25MG</u>	<u>A204279</u>	<u>002</u>	Aug 20, 2014
AB			<u>50MG</u>	<u>A204279</u>	<u>003</u>	Aug 20, 2014
AB		HETERO LABS LTD III	<u>10MG</u>	<u>A040805</u>	<u>001</u>	May 29, 2008
AB			<u>25MG</u>	<u>A040805</u>	<u>002</u>	May 29, 2008
AB			<u>50MG</u>	<u>A040805</u>	<u>003</u>	May 29, 2008
AB		INVAGEN PHARMS	<u>10MG</u>	<u>A040812</u>	<u>001</u>	Mar 12, 2008
AB			<u>25MG</u>	<u>A040812</u>	<u>002</u>	Mar 12, 2008
AB			<u>50MG</u>	<u>A040812</u>	<u>003</u>	Mar 12, 2008
AB		KVK TECH	<u>10MG</u>	<u>A040786</u>	<u>001</u>	Mar 20, 2007
AB			<u>25MG</u>	<u>A040786</u>	<u>002</u>	Mar 20, 2007
AB			<u>50MG</u>	<u>A040786</u>	<u>003</u>	Mar 20, 2007
AB		NORTHSTAR HLTHCARE	<u>10MG</u>	<u>A040840</u>	<u>002</u>	Mar 31, 2008
AB			<u>25MG</u>	<u>A040840</u>	<u>003</u>	Mar 31, 2008
AB			<u>50MG</u>	<u>A040840</u>	<u>001</u>	Mar 31, 2008
AB		NUVO PHARM	<u>10MG</u>	<u>A207120</u>	<u>001</u>	Mar 29, 2017
AB			<u>50MG</u>	<u>A207122</u>	<u>001</u>	Mar 29, 2017
AB		NUVO PHARMS INC	<u>25MG</u>	<u>A207121</u>	<u>001</u>	Mar 29, 2017
AB	!	PLIVA	<u>10MG</u>	<u>A088617</u>	<u>001</u>	Jan 10, 1986
AB	!		<u>25MG</u>	<u>A088618</u>	<u>001</u>	Jan 10, 1986
AB	!		<u>50MG</u>	<u>A088619</u>	<u>001</u>	Jan 10, 1986
AB		PRINSTON INC	<u>10MG</u>	<u>A040579</u>	<u>001</u>	May 27, 2005
AB			<u>25MG</u>	<u>A040574</u>	<u>001</u>	May 27, 2005
AB			<u>50MG</u>	<u>A040580</u>	<u>001</u>	May 27, 2005

HYDROXYZINE PAMOATE

CAPSULE; ORAL

HYDROXYZINE PAMOATE

AB		BARR	<u>EQ 25MG HYDROCHLORIDE</u>	<u>A088496</u>	<u>001</u>	Jun 15, 1984
AB			<u>EQ 50MG HYDROCHLORIDE</u>	<u>A088487</u>	<u>001</u>	Jun 15, 1984
AB		HERITAGE PHARMA	<u>EQ 25MG HYDROCHLORIDE</u>	<u>A201507</u>	<u>001</u>	Jun 03, 2013
AB			<u>EQ 50MG HYDROCHLORIDE</u>	<u>A201507</u>	<u>002</u>	Jun 03, 2013
AB		IMPAX LABS INC	<u>EQ 25MG HYDROCHLORIDE</u>	<u>A040156</u>	<u>001</u>	Jul 15, 1996
AB			<u>EQ 50MG HYDROCHLORIDE</u>	<u>A040156</u>	<u>002</u>	Jul 15, 1996
AB		SANDOZ	<u>EQ 25MG HYDROCHLORIDE</u>	<u>A087479</u>	<u>001</u>	
AB	!		<u>EQ 50MG HYDROCHLORIDE</u>	<u>A086183</u>	<u>001</u>	
		BARR	EQ 100MG HYDROCHLORIDE	A088488	001	Jun 15, 1984

IBANDRONATE SODIUM

INJECTABLE; INTRAVENOUS

BONIVA

AP	+	ROCHE	<u>EQ 3MG BASE/3ML</u>	<u>N021858</u>	<u>001</u>	Jan 06, 2006
AP		ACCORD HLTHCARE	<u>EQ 3MG BASE/3ML</u>	<u>A206058</u>	<u>001</u>	Feb 05, 2016
AP		APOTEX	<u>EQ 3MG BASE/3ML</u>	<u>A204222</u>	<u>001</u>	Oct 16, 2015
AP		AUROBINDO PHARMA	<u>EQ 3MG BASE/3ML</u>	<u>A205332</u>	<u>001</u>	Aug 19, 2015

PRESCRIPTION DRUG PRODUCT LIST

IBANDRONATE SODIUM

INJECTABLE; INTRAVENOUS

IBANDRONATE SODIUM

LTD

<u>AP</u>	MYLAN LABS LTD	<u>EQ 3MG BASE/3ML</u>	<u>A202671 001</u>	Sep 02, 2014
<u>AP</u>	SAGENT PHARMS INC	<u>EQ 3MG BASE/3ML</u>	<u>A202235 001</u>	Sep 02, 2014
<u>AP</u>	SUN PHARM	<u>EQ 3MG BASE/3ML</u>	<u>A090853 001</u>	Feb 14, 2014

TABLET; ORAL

BONIVA

<u>AB</u>	<u>+</u> !	HOFFMANN LA ROCHE	<u>EQ 150MG BASE</u>	<u>N021455 002</u>	Mar 24, 2005
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IBANDRONATE SODIUM

<u>AB</u>		APOTEX INC	<u>EQ 150MG BASE</u>	<u>A078948 001</u>	Mar 19, 2012
<u>AB</u>		AUROBINDO PHARMA LTD	<u>EQ 150MG BASE</u>	<u>A204502 001</u>	Mar 11, 2016
<u>AB</u>		DR REDDYS LABS LTD	<u>EQ 150MG BASE</u>	<u>A078997 001</u>	Apr 30, 2012
<u>AB</u>		MACLEODS PHARMS LTD	<u>EQ 150MG BASE</u>	<u>A206887 001</u>	Oct 31, 2017
<u>AB</u>		ORCHID HLTHCARE	<u>EQ 150MG BASE</u>	<u>A078998 001</u>	Mar 19, 2012
<u>AB</u>		WATSON LABS TEVA	<u>EQ 150MG BASE</u>	<u>A079003 001</u>	Mar 20, 2012

IBRUTINIB

CAPSULE; ORAL

IMBRUVICA

<u>+</u>		PHARMACYCLICS INC	70MG	N205552 002	Dec 20, 2017
<u>+</u> !			140MG	N205552 001	Nov 13, 2013

TABLET; ORAL

IMBRUVICA

<u>+</u>		PHARMACYCLICS INC	140MG	N210563 001	Feb 16, 2018
<u>+</u>			280MG	N210563 002	Feb 16, 2018
<u>+</u>			420MG	N210563 003	Feb 16, 2018
<u>+</u> !			560MG	N210563 004	Feb 16, 2018

IBUPROFEN

SOLUTION; INTRAVENOUS

CALDOLOR

<u>+</u> !		CUMBERLAND PHARMS	800MG/8ML (100MG/ML)	N022348 002	Jun 11, 2009
<u>+</u> !			800MG/200ML (4MG/ML)	N022348 003	Jan 25, 2019

SUSPENSION; ORAL

IBUPROFEN

<u>AB</u>	<u>!</u>	ACTAVIS MID ATLANTIC	<u>100MG/5ML</u>	<u>A074978 001</u>	Mar 25, 1998
<u>AB</u>		HI TECH	<u>100MG/5ML</u>	<u>A205647 001</u>	Nov 03, 2016
<u>AB</u>		L PERRIGO CO	<u>100MG/5ML</u>	<u>A076925 001</u>	Sep 23, 2004
<u>AB</u>		TARO	<u>100MG/5ML</u>	<u>A209204 001</u>	Jun 23, 2017
		AUROBINDO PHARMA LTD	100MG/5ML	A209178 001	Feb 16, 2018

TABLET; ORAL

IBU-TAB

<u>AB</u>		ALRA	<u>400MG</u>	<u>A071058 001</u>	Aug 11, 1988
<u>AB</u>			<u>600MG</u>	<u>A071059 001</u>	Aug 11, 1988

IBUPROFEN

<u>AB</u>		AMNEAL PHARMS NY	<u>400MG</u>	<u>A071334 001</u>	Nov 25, 1986
<u>AB</u>			<u>400MG</u>	<u>A078558 001</u>	Jun 18, 2007
<u>AB</u>			<u>600MG</u>	<u>A071335 001</u>	Nov 25, 1986
<u>AB</u>			<u>600MG</u>	<u>A078558 002</u>	Jun 18, 2007
<u>AB</u>			<u>800MG</u>	<u>A071935 001</u>	Oct 13, 1987
<u>AB</u>			<u>800MG</u>	<u>A078558 003</u>	Jun 18, 2007
<u>AB</u>		CONTRACT PHARMACAL	<u>400MG</u>	<u>A071268 002</u>	Oct 15, 1986
<u>AB</u>			<u>600MG</u>	<u>A071268 001</u>	Oct 15, 1986
<u>AB</u>			<u>800MG</u>	<u>A071268 003</u>	Jul 01, 1988
<u>AB</u>		DR REDDYS LA	<u>400MG</u>	<u>A075682 001</u>	Nov 14, 2001
<u>AB</u>			<u>600MG</u>	<u>A075682 002</u>	Nov 14, 2001
<u>AB</u>	<u>!</u>		<u>800MG</u>	<u>A075682 003</u>	Nov 14, 2001
<u>AB</u>		DR REDDYS LABS INC	<u>400MG</u>	<u>A076112 001</u>	Oct 31, 2001
<u>AB</u>			<u>600MG</u>	<u>A076112 002</u>	Oct 31, 2001
<u>AB</u>			<u>800MG</u>	<u>A076112 003</u>	Oct 31, 2001
<u>AB</u>		GRANULES INDIA LTD	<u>400MG</u>	<u>A091625 001</u>	Sep 15, 2015
<u>AB</u>			<u>600MG</u>	<u>A091625 002</u>	Sep 15, 2015
<u>AB</u>			<u>800MG</u>	<u>A091625 003</u>	Sep 15, 2015
<u>AB</u>		MARKSANS PHARMA	<u>400MG</u>	<u>A090796 001</u>	Dec 21, 2010
<u>AB</u>			<u>600MG</u>	<u>A090796 002</u>	Dec 21, 2010
<u>AB</u>			<u>800MG</u>	<u>A090796 003</u>	Dec 21, 2010
<u>AB</u>		PERRIGO PHARMS CO	<u>400MG</u>	<u>A077114 001</u>	Jul 18, 2005
<u>AB</u>			<u>600MG</u>	<u>A077114 002</u>	Jul 18, 2005
<u>AB</u>			<u>800MG</u>	<u>A077114 003</u>	Jul 18, 2005

PRESCRIPTION DRUG PRODUCT LIST

IBUPROFEN

TABLET; ORAL

IBUPROFEN

<u>AB</u>	SHANDONG XINHUA	<u>400MG</u>	<u>A202413 001</u>	Nov 23, 2016
<u>AB</u>		<u>600MG</u>	<u>A202413 002</u>	Nov 23, 2016
<u>AB</u>		<u>800MG</u>	<u>A202413 003</u>	Nov 23, 2016
<u>AB</u>	STRIDES PHARMA	<u>400MG</u>	<u>A078329 001</u>	Feb 05, 2009
<u>AB</u>		<u>600MG</u>	<u>A078329 002</u>	Feb 05, 2009
<u>AB</u>		<u>800MG</u>	<u>A078329 003</u>	Feb 05, 2009
<u>AB</u>	VINTAGE PHARMS	<u>400MG</u>	<u>A071644 001</u>	Feb 01, 1988

IBUPROFEN LYISINE

INJECTABLE; INTRAVENOUS

IBUPROFEN LYISINE

<u>AP</u>	XGEN PHARMS	<u>EQ 20MG BASE/2ML (EQ 10MG BASE/ML)</u>	<u>A202402 001</u>	Mar 30, 2016
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NEOPROFEN

<u>AP</u>	+! RECORDATI RARE	<u>EQ 20MG BASE/2ML (EQ 10MG BASE/ML)</u>	<u>N021903 001</u>	Apr 13, 2006
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IBUPROFEN; OXYCODONE HYDROCHLORIDE

TABLET; ORAL

OXYCODONE HYDROCHLORIDE AND IBUPROFEN

<u>AB</u>	ACTAVIS ELIZABETH	<u>400MG;5MG</u>	<u>A078769 001</u>	Jan 04, 2008
<u>AB</u>	! BARR LABS INC	<u>400MG;5MG</u>	<u>A078316 001</u>	Nov 29, 2007

IBUTILIDE FUMARATE

INJECTABLE; INJECTION

CORVERT

<u>AP</u>	+! PHARMACIA AND UPJOHN	<u>0.1MG/ML</u>	<u>N020491 001</u>	Dec 28, 1995
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IBUTILIDE FUMARATE

<u>AP</u>	MYLAN INSTITUTIONAL	<u>0.1MG/ML</u>	<u>A090643 001</u>	Jan 11, 2010
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ICATIBANT ACETATE

INJECTABLE; SUBCUTANEOUS

FIRAZYR

<u>AP</u>	+! SHIRE ORPHAN THERAP	<u>EQ 30MG BASE/3ML (EQ 10MG BASE/ML)</u>	<u>N022150 001</u>	Aug 25, 2011
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ICATIBANT ACETATE

<u>AP</u>	JIANGSU HANSON PHARM	<u>EQ 30MG BASE/3ML (EQ 10MG BASE/ML)</u>	<u>A211021 001</u>	Mar 09, 2020
<u>AP</u>	TEVA PHARMS USA	<u>EQ 30MG BASE/3ML (EQ 10MG BASE/ML)</u>	<u>A210118 001</u>	Jul 15, 2019

ICODEXTRIN

SOLUTION; INTRAPERITONEAL

EXTRANEAL

+!	BAXTER HLTHCARE	7.5GM/100ML	N021321 001	Dec 20, 2002
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ICOSAPENT ETHYL

CAPSULE; ORAL

VASCEPA

+	AMARIN PHARMS	500MG	N202057 002	Feb 16, 2017
+!		1GM	N202057 001	Jul 26, 2012

IDARUBICIN HYDROCHLORIDE

INJECTABLE; INJECTION

IDAMYCIN PFS

<u>AP</u>	+! PHARMACIA AND UPJOHN	<u>1MG/ML</u>	<u>N050734 001</u>	Feb 17, 1997
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IDARUBICIN HYDROCHLORIDE

<u>AP</u>	FRESENIUS KABI USA	<u>1MG/ML</u>	<u>A065440 001</u>	Aug 04, 2009
<u>AP</u>	WEST-WARD PHARMS INT	<u>1MG/ML</u>	<u>A065275 001</u>	Dec 14, 2006

<u>AP</u>		<u>1MG/ML</u>	<u>A065288 001</u>	May 15, 2007
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IDARUBICIN HYDROCHLORIDE PFS

<u>AP</u>	TEVA PHARMS USA	<u>1MG/ML</u>	<u>A065036 001</u>	May 01, 2002
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IDELALISIB

TABLET; ORAL

ZYDELIG

+	GILEAD SCIENCES INC	100MG	N205858 001	Jul 23, 2014
+!		150MG	N205858 002	Jul 23, 2014

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

PRESCRIPTION DRUG PRODUCT LIST

IFOSFAMIDE

INJECTABLE; INJECTION

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>	!	TEVA PHARMS USA	<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
<u>AP</u>		WEST-WARD PHARMS INT	<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

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; INHALATION

VENTAVIS

+	!	ACTELION PHARMS LTD	10MCG/ML (10MCG/ML)	N021779 002	Dec 08, 2005
+	!		20MCG/ML (20MCG/ML)	N021779 003	Aug 07, 2009

IMATINIB MESYLATE

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>		BRECKENRIDGE	<u>EQ 100MG BASE</u>	<u>A205990 001</u>	Feb 08, 2019
<u>AB</u>			<u>EQ 400MG BASE</u>	<u>A205990 002</u>	Feb 08, 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>		HIKMA	<u>EQ 100MG BASE</u>	<u>A207586 001</u>	Jul 13, 2018
<u>AB</u>			<u>EQ 400MG BASE</u>	<u>A207586 002</u>	Jul 13, 2018
<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>		SHILPA MEDICARE LTD	<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>		WOCKHARDT BIO AG	<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

IMIGLUCERASE

INJECTABLE; INJECTION

CEREZYME

+		GENZYME	200 UNITS/VIAL	N020367 001	May 23, 1994
+	!		400 UNITS/VIAL	N020367 002	Sep 22, 1999

IMIPRAMINE HYDROCHLORIDE

TABLET; ORAL

IMIPRAMINE HYDROCHLORIDE

<u>AB</u>		LEADING PHARMA LLC	<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>		PAR PHARM	<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>		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>		SPEGX LLC	<u>10MG</u>	<u>A087846 002</u>	May 22, 1984
<u>AB</u>			<u>25MG</u>	<u>A087846 003</u>	May 22, 1984

PRESCRIPTION DRUG PRODUCT LIST

IMIPRAMINE HYDROCHLORIDE

TABLET; ORAL

IMIPRAMINE HYDROCHLORIDE

<u>AB</u>	!	SUN PHARM INDUSTRIES	<u>10MG</u>	<u>A081048 001</u>	Jun 05, 1990
<u>AB</u>			<u>25MG</u>	<u>A081049 001</u>	Jun 05, 1990
<u>AB</u>			<u>50MG</u>	<u>A081050 001</u>	Jun 05, 1990
<u>TOFRANIL</u>					
<u>AB</u>	!	SPECGX LLC	<u>50MG</u>	<u>A087846 001</u>	May 22, 1984

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

ALDARA

<u>AB</u>	+	!	BAUSCH	<u>5%</u>	<u>N020723 001</u>	Feb 27, 1997
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IMIQUIMOD

<u>AB</u>			ANDA REPOSITORY	<u>5%</u>	<u>A091044 001</u>	Feb 28, 2011
<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 GENERICS	<u>5%</u>	<u>A201994 001</u>	Mar 06, 2012
<u>AB</u>			PERRIGO ISRAEL	<u>5%</u>	<u>A078837 001</u>	Sep 07, 2010
<u>AB</u>			TARO	<u>5%</u>	<u>A200173 001</u>	Apr 15, 2011
<u>ZYCLARA</u>						
	+	!	BAUSCH	2.5%	N022483 002	Jul 15, 2011
	+	!		3.75%	N022483 001	Mar 25, 2010

INAMRINONE LACTATE

INJECTABLE; INJECTION

AMRINONE LACTATE

!	WEST-WARD PHARMS INT	EQ 5MG BASE/ML	<u>A075513 001</u>	May 09, 2000
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INDACATEROL MALEATE

POWDER; INHALATION

ARCAPTA NEOHALER

+	!	SUNOVION PHARMS INC	EQ 75MCG BASE	<u>N022383 001</u>	Jul 01, 2011
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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 INC	<u>1.25MG</u>	<u>A074299 002</u>	Apr 29, 1996
<u>AB</u>			<u>1.25MG</u>	<u>A075201 001</u>	Dec 04, 1998
<u>AB</u>			<u>2.5MG</u>	<u>A074299 001</u>	Jul 27, 1995
<u>AB</u>			<u>2.5MG</u>	<u>A075201 002</u>	Dec 04, 1998
<u>AB</u>		MYLAN	<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

INDINAVIR SULFATE

CAPSULE; ORAL

CRIXIVAN

+	MERCK SHARP DOHME	EQ 200MG BASE	<u>N020685 003</u>	Mar 13, 1996
+	!		<u>N020685 001</u>	Mar 13, 1996

INDIUM IN-111 CHLORIDE

INJECTABLE; INJECTION

INDIUM IN 111 CHLORIDE

+	!	CURIUM	5mCi/0.5ML	<u>N019841 001</u>	Sep 27, 1994
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PRESCRIPTION DRUG PRODUCT LIST

INDIUM IN-111 OXYQUINOLINE

INJECTABLE; INJECTION

INDIUM IN 111 OXYQUINOLINE

<u>AP</u>	BWXT ITG	<u>1mCi/ML</u>	<u>A202586</u>	<u>001</u>	Jul 25, 2018
<u>AP</u>	+! GE HEALTHCARE	<u>1mCi/ML</u>	<u>N019044</u>	<u>001</u>	Dec 24, 1985

INDIUM IN-111 PENTETATE DISODIUM

INJECTABLE; INTRATHECAL

MPI INDIUM DTPA IN 111

+!	GE HEALTHCARE	1mCi/ML	N017707	001	Feb 18, 1982
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INDIUM IN-111 PENTETREOTIDE KIT

INJECTABLE; INJECTION

OCTREOSCAN

+!	CURIUM	3mCi/ML	N020314	001	Jun 02, 1994
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INDOCYANINE GREEN

INJECTABLE; INJECTION

IC-GREEN

<u>AP</u>	+! AKORN	<u>25MG/VIAL</u>	<u>N011525</u>	<u>001</u>	
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INDOCYANINE GREEN

<u>AP</u>	DIAGNOSTIC GREEN	<u>25MG/VIAL</u>	<u>A040811</u>	<u>001</u>	Nov 21, 2007
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POWDER; INTRAVENOUS, INTERSTITIAL

SPY AGENT GREEN KIT

+!	NOVADAQ TECH	25MG/VIAL	N211580	001	Nov 21, 2018
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INDOMETHACIN

CAPSULE; ORAL

INDOMETHACIN

<u>AB</u>	CADILA	<u>25MG</u>	<u>A090403</u>	<u>001</u>	Nov 15, 2010
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<u>AB</u>		<u>50MG</u>	<u>A090403</u>	<u>002</u>	Nov 15, 2010
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<u>AB</u>	GLENMARK GENERICS	<u>25MG</u>	<u>A091276</u>	<u>001</u>	Dec 22, 2010
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<u>AB</u>	!	<u>50MG</u>	<u>A091276</u>	<u>002</u>	Dec 22, 2010
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<u>AB</u>	HERITAGE	<u>25MG</u>	<u>N018851</u>	<u>001</u>	May 18, 1984
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<u>AB</u>		<u>50MG</u>	<u>N018851</u>	<u>002</u>	May 18, 1984
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<u>AB</u>	HERITAGE PHARMA	<u>25MG</u>	<u>A070719</u>	<u>001</u>	Feb 12, 1986
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<u>AB</u>		<u>50MG</u>	<u>A070756</u>	<u>001</u>	Feb 12, 1986
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<u>AB</u>	HETERO LABS LTD III	<u>25MG</u>	<u>A091240</u>	<u>001</u>	Apr 12, 2011
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<u>AB</u>		<u>50MG</u>	<u>A091240</u>	<u>002</u>	Apr 12, 2011
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<u>AB</u>	JUBILANT GENERICS	<u>25MG</u>	<u>A205215</u>	<u>001</u>	Aug 25, 2017
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<u>AB</u>		<u>50MG</u>	<u>A205215</u>	<u>002</u>	Aug 25, 2017
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<u>AB</u>	SANDOZ	<u>25MG</u>	<u>A070673</u>	<u>001</u>	Apr 29, 1987
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<u>AB</u>		<u>50MG</u>	<u>A070674</u>	<u>001</u>	Apr 29, 1987
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TIVORBEX

+	GENUS	20MG	N204768	001	Feb 24, 2014
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+!		40MG	N204768	002	Feb 24, 2014
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CAPSULE, EXTENDED RELEASE; ORAL

INDOMETHACIN

<u>AB</u>	AMNEAL PHARMS	<u>75MG</u>	<u>A091549</u>	<u>001</u>	Dec 01, 2010
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<u>AB</u>	AUROBINDO PHARMA LTD	<u>75MG</u>	<u>A204243</u>	<u>001</u>	Dec 27, 2016
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<u>AB</u>	AVANTHI INC	<u>75MG</u>	<u>A079175</u>	<u>001</u>	Mar 06, 2009
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<u>AB</u>	CHARTWELL RX	<u>75MG</u>	<u>A200529</u>	<u>001</u>	Nov 30, 2010
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<u>AB</u>	GLENMARK PHARMS LTD	<u>75MG</u>	<u>A203501</u>	<u>001</u>	Jun 22, 2017
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<u>AB</u>	HETERO LABS LTD III	<u>75MG</u>	<u>A201807</u>	<u>001</u>	Sep 28, 2012
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<u>AB</u>	JUBILANT GENERICS	<u>75MG</u>	<u>A202706</u>	<u>001</u>	Oct 05, 2015
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<u>AB</u>	MYLAN	<u>75MG</u>	<u>A202139</u>	<u>001</u>	Mar 20, 2014
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<u>AB</u>	NOVAST LABS	<u>75MG</u>	<u>A204853</u>	<u>001</u>	May 08, 2017
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<u>AB</u>	!	<u>75MG</u>	<u>A074464</u>	<u>001</u>	May 28, 1998
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<u>AB</u>	ZYDUS PHARMS	<u>75MG</u>	<u>A202711</u>	<u>001</u>	Sep 25, 2017
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INJECTABLE; INJECTION

INDOMETHACIN

+!	FRESENIUS KABI USA	EQ 1MG BASE/VIAL	N022536	001	Mar 17, 2010
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SUPPOSITORY; RECTAL

INDOMETHACIN

!	ACP NIMBLE	50MG	A073314	001	Aug 31, 1992
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SUSPENSION; ORAL

INDOCIN

+!	EGALET	25MG/5ML	N018332	001	Oct 10, 1985
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PRESCRIPTION DRUG PRODUCT LISTINDOMETHACIN SODIUM

INJECTABLE; INJECTION

INDOCIN

AP	+ !	RECORDATI RARE	<u>EQ 1MG BASE/VIAL</u>	<u>N018878 001</u>	Jan 30, 1985
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INDOMETHACIN SODIUM

AP		HOSPIRA INC	<u>EQ 1MG BASE/VIAL</u>	<u>A204118 001</u>	Apr 19, 2016
AP		NAVINTA LLC	<u>EQ 1MG BASE/VIAL</u>	<u>A206561 001</u>	Jul 19, 2017
AP		WEST-WARD PHARMS INT	<u>EQ 1MG BASE/VIAL</u>	<u>A078713 001</u>	Jul 16, 2008

INGENOL MEBUTATE

GEL; TOPICAL

INGENOL MEBUTATE

AB		PERRIGO UK FINCO	<u>0.015%</u>	<u>A209018 001</u>	Jan 07, 2019
AB			<u>0.05%</u>	<u>A209019 001</u>	Jan 09, 2019
AB	+ !	LEO LABS	<u>0.015%</u>	<u>N202833 001</u>	Jan 23, 2012
AB	+ !		<u>0.05%</u>	<u>N202833 002</u>	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
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INSULIN ASPART

SOLUTION; INTRAVENOUS, SUBCUTANEOUS

FIASP

+ !	NOVO	1000 UNITS/10ML (100 UNITS/ML)	N208751 001	Sep 29, 2017
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SOLUTION; SUBCUTANEOUS

FIASP FLEXTOUCH

+ !	NOVO	300 UNITS/3ML (100 UNITS/ML)	N208751 002	Sep 29, 2017
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FIASP PENFILL

+ !	NOVO	300 UNITS/3ML (100 UNITS/ML)	N208751 003	Sep 24, 2018
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INSULIN ASPART PROTAMINE RECOMBINANT; INSULIN ASPART RECOMBINANT

INJECTABLE; SUBCUTANEOUS

NOVOLOG MIX 70/30

+ !	NOVO NORDISK INC	700 UNITS/10ML; 300 UNITS/10ML (70 UNITS/ML; 30 UNITS/ML)	N021172 001	Nov 01, 2001
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NOVOLOG MIX 70/30 FLEXPEN

+ !	NOVO NORDISK INC	210 UNITS/3ML; 90 UNITS/3ML (70 UNITS/ML; 30 UNITS/ML)	N021172 004	May 03, 2002
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INSULIN ASPART RECOMBINANT

INJECTABLE; SUBCUTANEOUS

NOVOLOG

+ !	NOVO NORDISK INC	1000 UNITS/10ML (100 UNITS/ML)	N020986 001	Jun 07, 2000
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NOVOLOG FLEXPEN

+ !	NOVO NORDISK INC	300 UNITS/3ML (100 UNITS/ML)	N020986 003	Jan 19, 2001
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NOVOLOG PENFILL

+ !	NOVO NORDISK INC	300 UNITS/3ML (100 UNITS/ML)	N020986 002	Jun 07, 2000
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INSULIN DEGLUDEC

SOLUTION; SUBCUTANEOUS

TRESIBA

+	NOVO	300 UNITS/3ML (100 UNITS/ML)	N203314 001	Sep 25, 2015
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+ !		600 UNITS/3ML (200 UNITS/ML)	N203314 002	Sep 25, 2015
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		1000 UNITS/10ML (100 UNITS/ML)	N203314 003	Nov 21, 2018
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INSULIN DEGLUDEC; LIRAGLUTIDE

SOLUTION; SUBCUTANEOUS

XULTOPHY 100/3.6

+ !	NOVO	300 UNITS/3ML; 10.8MG/3ML (100 UNITS/ML; 3.6MG/ML)	N208583 001	Nov 21, 2016
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INSULIN DETEMIR RECOMBINANT

INJECTABLE; SUBCUTANEOUS

LEVEMIR

+ !	NOVO NORDISK INC	1000 UNITS/10ML (100 UNITS/ML)	N021536 001	Jun 16, 2005
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LEVEMIR FLEXTOUCH

+ !	NOVO NORDISK INC	300 UNITS/3ML (100 UNITS/ML)	N021536 005	Oct 31, 2013
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PRESCRIPTION DRUG PRODUCT LISTINSULIN GLARGINE

SOLUTION;SUBCUTANEOUS

BASAGLAR

ELI LILLY AND CO 300 UNITS/3ML (100 UNITS/ML) N205692 001 Dec 16, 2015

INSULIN GLARGINE RECOMBINANT

INJECTABLE;INJECTION

LANTUS

+! SANOFI AVENTIS US 100 UNITS/ML N021081 001 Apr 20, 2000

LANTUS SOLOSTAR

+! SANOFI AVENTIS US 300 UNITS/3ML (100 UNITS/ML) N021081 002 Apr 27, 2007

SOLUTION;SUBCUTANEOUS

TOUJEO MAX SOLOSTAR

+! SANOFI US SERVICES 900 UNITS/3ML (300 UNITS/ML) N206538 002 Mar 26, 2018

TOUJEO SOLOSTAR

+! SANOFI US SERVICES 450 UNITS/1.5ML (300 UNITS/ML) N206538 001 Feb 25, 2015

INSULIN GLARGINE; LIXISENATIDE

SOLUTION;SUBCUTANEOUS

SOLIQUA 100/33

+! SANOFI-AVENTIS US 300 UNITS/3ML;99MCG/3ML (100 UNITS/ML;33MCG/ML) N208673 001 Nov 21, 2016

INSULIN GLULISINE RECOMBINANT

INJECTABLE;INTRAVENOUS, SUBCUTANEOUS

APIDRA

+! SANOFI AVENTIS US 1000 UNITS/10ML (100 UNITS/ML) N021629 001 Apr 16, 2004

INJECTABLE;SUBCUTANEOUS

APIDRA SOLOSTAR

+ SANOFI AVENTIS US 300 UNITS/3ML N021629 003 Feb 24, 2009

INSULIN HUMAN

SOLUTION;INTRAVENOUS

MYXREDLIN

+! BAXTER HLTHCARE 100 UNITS/100ML (1 UNIT/ML) N208157 001 Jun 20, 2019
CORP

SOLUTION;SUBCUTANEOUS

HUMULIN R

+! LILLY 10000 UNITS/20ML (500 UNITS/ML) N018780 004 Mar 31, 1994

HUMULIN R KWIKPEN

+! LILLY 1500 UNITS/3ML (500 UNITS/ML) N018780 002 Dec 29, 2015

INSULIN LISPRO

SOLUTION;INTRAVENOUS, SUBCUTANEOUS

ADMELOG

+ SANOFI-AVENTIS US 300 UNITS/3ML (100 UNITS/ML) N209196 003 Oct 19, 2018

+ 1000 UNITS/10ML (100 UNITS/ML) N209196 001 Dec 11, 2017

ADMELOG SOLOSTAR

+ SANOFI-AVENTIS US 300 UNITS/3ML (100 UNITS/ML) N209196 002 Dec 11, 2017

INSULIN LISPRO PROTAMINE RECOMBINANT; INSULIN LISPRO RECOMBINANT

INJECTABLE;INJECTION

HUMALOG MIX 50/50

+! LILLY 50 UNITS/ML;50 UNITS/ML N021018 001 Dec 22, 1999

HUMALOG MIX 50/50 KWIKPEN

+! LILLY 50 UNITS/ML;50 UNITS/ML N021018 002 Sep 06, 2007

HUMALOG MIX 75/25

+! LILLY 75 UNITS/ML;25 UNITS/ML N021017 001 Dec 22, 1999

HUMALOG MIX 75/25 KWIKPEN

+! LILLY 75 UNITS/ML;25 UNITS/ML N021017 002 Sep 06, 2007

INSULIN LISPRO RECOMBINANT

INJECTABLE;INJECTION

HUMALOG

+! LILLY 100 UNITS/ML N020563 001 Jun 14, 1996

HUMALOG KWIKPEN

+! LILLY 100 UNITS/ML N020563 003 Sep 06, 2007

+! 200UNITS/ML N020563 004 Jan 06, 2017

HUMALOG TEMPO PEN

+! LILLY 100UNITS/ML N020563 005 Nov 15, 2019

SOLUTION;SUBCUTANEOUS

HUMALOG KWIKPEN

+! ELI LILLY AND CO 200 UNITS/ML N205747 001 May 26, 2015

PRESCRIPTION DRUG PRODUCT LISTINSULIN RECOMBINANT HUMAN

POWDER; INHALATION

AFREZZA

+	MANNKIND	4 UNITS/INH	N022472 001	Jun 27, 2014
+	!	8 UNITS/INH	N022472 002	Jun 27, 2014
+		12 UNITS/INH	N022472 003	Apr 17, 2015

IOBENGUANE I-131

SOLUTION; INTRAVENOUS

AZEDRA

+	!	PROGENICS PHARMS INC	15mCi/ML	N209607 001	Jul 30, 2018
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IOBENGUANE SULFATE I-123

SOLUTION; INTRAVENOUS

ADREVIEW

+	!	GE HEALTHCARE	10mCi/5ML (2mCi/ML)	N022290 001	Sep 19, 2008
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IODIXANOL

INJECTABLE; INJECTION

VISIPAQUE 270

+	!	GE HEALTHCARE	55%	N020351 001	Mar 22, 1996
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VISIPAQUE 320

+	!	GE HEALTHCARE	65.2%	N020351 002	Mar 22, 1996
			65.2%	N020808 002	Aug 29, 1997

IOFLUPANE I-123

SOLUTION; INTRAVENOUS

DATSCAN

+	!	GE HLTHCARE INC	5mCi/2.5ML (2mCi/ML)	N022454 001	Jan 14, 2011
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IOHEXOL

FOR SOLUTION; ORAL

ORALTAG

INTERPHARMA PRAHA
AS

			9.7GM/BOT	N205383 001	Mar 26, 2015
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INJECTABLE; INJECTION

OMNIPAQUE 140

+	!	GE HEALTHCARE	30.2%	N018956 005	Nov 30, 1988
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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
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OMNIPAQUE 240

+	!	GE HEALTHCARE	51.8%	N018956 002	Dec 26, 1985
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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
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OMNIPAQUE 9

+	!	GE HEALTHCARE	1.9%	N018956 008	Apr 17, 2018
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IOPAMIDOL

INJECTABLE; INJECTION

ISOVUE-300

AP	+	!	BRACCO	61%	N018735 002	Dec 31, 1985
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ISOVUE-370

AP	+	!	BRACCO	76%	N018735 003	Dec 31, 1985
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SCANLUX-300

AP			SANOCHEMIA CORP USA	61%	A090394 001	Jun 18, 2010
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SCANLUX-370

AP			SANOCHEMIA CORP USA	76%	A090394 002	Jun 18, 2010
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ISOVUE-200

+	!	BRACCO	41%	N018735 006	Jul 07, 1987
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ISOVUE-250

+	!	BRACCO	51%	N018735 007	Jul 06, 1992
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+	!		51%	N020327 002	Oct 12, 1994
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ISOVUE-300

+	!	BRACCO	61%	N020327 003	Oct 12, 1994
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PRESCRIPTION DRUG PRODUCT LIST

IOPAMIDOL

INJECTABLE; INJECTION

ISOVUE-370

+! BRACCO 76% N020327 004 Oct 12, 1994

ISOVUE-M 200

+! BRACCO 41% N018735 001 Dec 31, 1985

ISOVUE-M 300

+! BRACCO 61% N018735 004 Dec 31, 1985

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 240

+! LIEBEL-FLARSHEIM 51% N019710 002 Dec 30, 1988

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

SOLUTION; INHALATION

IPRATROPIUM BROMIDE

AN AUROBINDO PHARMA LTD **0.02%** **A206543 001** Oct 27, 2016

AN LANDELA PHARM **0.02%** **A077072 001** Jul 19, 2005

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

AN WATSON LABS **0.02%** **A076291 001** May 09, 2005

SPRAY, METERED; NASAL

IPRATROPIUM BROMIDE

AB APOTEX INC **0.021MG/SPRAY** **A076156 001** Apr 18, 2003

AB **0.042MG/SPRAY** **A076155 001** Apr 18, 2003

AB BAUSCH AND LOMB **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

IRBESARTAN

TABLET; ORAL

AVAPRO

AB + SANOFI AVENTIS US **75MG** **N020757 001** Sep 30, 1997

AB + **150MG** **N020757 002** Sep 30, 1997

AB +! **300MG** **N020757 003** Sep 30, 1997

PRESCRIPTION DRUG PRODUCT LIST

3-241 (of 453)

IRBESARTAN

TABLET; ORAL

IRBESARTAN

<u>AB</u>	ALEMbic PHARMS LTD	<u>75MG</u>	<u>A091236 001</u>	Oct 15, 2012
<u>AB</u>		<u>150MG</u>	<u>A091236 002</u>	Oct 15, 2012
<u>AB</u>		<u>300MG</u>	<u>A091236 003</u>	Oct 15, 2012
<u>AB</u>	AMNEAL PHARMS	<u>75MG</u>	<u>A204740 001</u>	Apr 17, 2018
<u>AB</u>		<u>150MG</u>	<u>A204740 002</u>	Apr 17, 2018
<u>AB</u>		<u>300MG</u>	<u>A204740 003</u>	Apr 17, 2018
<u>AB</u>	AUROBINDO PHARMA LTD	<u>75MG</u>	<u>A203081 001</u>	Sep 27, 2012
<u>AB</u>		<u>150MG</u>	<u>A203081 002</u>	Sep 27, 2012
<u>AB</u>		<u>300MG</u>	<u>A203081 003</u>	Sep 27, 2012
<u>AB</u>	CHARTWELL MOLECULAR	<u>75MG</u>	<u>A077205 001</u>	Nov 14, 2012
<u>AB</u>		<u>150MG</u>	<u>A077205 002</u>	Nov 14, 2012
<u>AB</u>		<u>300MG</u>	<u>A077205 003</u>	Nov 14, 2012
<u>AB</u>	HETERO LABS LTD V	<u>75MG</u>	<u>A202910 001</u>	Sep 27, 2012
<u>AB</u>		<u>150MG</u>	<u>A202910 002</u>	Sep 27, 2012
<u>AB</u>		<u>300MG</u>	<u>A202910 003</u>	Sep 27, 2012
<u>AB</u>	HISUN PHARM HANGZHOU	<u>75MG</u>	<u>A206194 001</u>	Jun 14, 2016
<u>AB</u>		<u>150MG</u>	<u>A206194 002</u>	Jun 14, 2016
<u>AB</u>		<u>300MG</u>	<u>A206194 003</u>	Jun 14, 2016
<u>AB</u>	JUBILANT GENERICS	<u>75MG</u>	<u>A203534 001</u>	Feb 23, 2015
<u>AB</u>		<u>150MG</u>	<u>A203534 002</u>	Feb 23, 2015
<u>AB</u>		<u>300MG</u>	<u>A203534 003</u>	Feb 23, 2015
<u>AB</u>	LUPIN LTD	<u>75MG</u>	<u>A201531 001</u>	Oct 15, 2012
<u>AB</u>		<u>150MG</u>	<u>A201531 002</u>	Oct 15, 2012
<u>AB</u>		<u>300MG</u>	<u>A201531 003</u>	Oct 15, 2012
<u>AB</u>	MACLEODS PHARMS LTD	<u>75MG</u>	<u>A202254 001</u>	Oct 03, 2012
<u>AB</u>		<u>150MG</u>	<u>A202254 002</u>	Oct 03, 2012
<u>AB</u>		<u>300MG</u>	<u>A202254 003</u>	Oct 03, 2012
<u>AB</u>	NEOPHARMA	<u>75MG</u>	<u>A203161 001</u>	Sep 27, 2012
<u>AB</u>		<u>150MG</u>	<u>A203161 002</u>	Sep 27, 2012
<u>AB</u>		<u>300MG</u>	<u>A203161 003</u>	Sep 27, 2012
<u>AB</u>	PRINSTON INC	<u>75MG</u>	<u>A203071 001</u>	Sep 27, 2012
<u>AB</u>		<u>150MG</u>	<u>A203071 002</u>	Sep 27, 2012
<u>AB</u>		<u>300MG</u>	<u>A203071 003</u>	Sep 27, 2012
<u>AB</u>	SANDOZ	<u>75MG</u>	<u>A077466 001</u>	Sep 27, 2012
<u>AB</u>		<u>150MG</u>	<u>A077466 002</u>	Sep 27, 2012
<u>AB</u>		<u>300MG</u>	<u>A077466 003</u>	Sep 27, 2012
<u>AB</u>	SCIEGEN PHARMS INC	<u>75MG</u>	<u>A204774 001</u>	Dec 07, 2015
<u>AB</u>		<u>150MG</u>	<u>A204774 002</u>	Dec 07, 2015
<u>AB</u>		<u>300MG</u>	<u>A204774 003</u>	Dec 07, 2015
<u>AB</u>	TEVA PHARMS	<u>75MG</u>	<u>A077159 001</u>	Mar 30, 2012
<u>AB</u>		<u>150MG</u>	<u>A077159 002</u>	Mar 30, 2012
<u>AB</u>		<u>300MG</u>	<u>A077159 003</u>	Mar 30, 2012
<u>AB</u>	UNICHEM LABS LTD	<u>75MG</u>	<u>A203020 001</u>	Dec 07, 2015
<u>AB</u>		<u>150MG</u>	<u>A203020 002</u>	Dec 07, 2015
<u>AB</u>		<u>300MG</u>	<u>A203020 003</u>	Dec 07, 2015
<u>AB</u>	ZYDUS PHARMS USA INC	<u>75MG</u>	<u>A079213 001</u>	Sep 27, 2012
<u>AB</u>		<u>150MG</u>	<u>A079213 002</u>	Sep 27, 2012
<u>AB</u>		<u>300MG</u>	<u>A079213 003</u>	Sep 27, 2012

IRINOTECAN HYDROCHLORIDE

INJECTABLE; INJECTION

CAMPTOSAR

<u>AP</u>	+	PFIZER INC	<u>40MG/2ML (20MG/ML)</u>	<u>N020571 001</u>	Jun 14, 1996
<u>AP</u>	+		<u>100MG/5ML (20MG/ML)</u>	<u>N020571 002</u>	Jun 14, 1996
<u>AP</u>	+		<u>300MG/15ML (20MG/ML)</u>	<u>N020571 003</u>	Aug 05, 2010

IRINOTECAN HYDROCHLORIDE

<u>AP</u>		ACCORD HLTHCARE	<u>40MG/2ML (20MG/ML)</u>	<u>A079068 001</u>	Nov 21, 2008
<u>AP</u>			<u>100MG/5ML (20MG/ML)</u>	<u>A079068 002</u>	Nov 21, 2008
<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>		AKORN	<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>		CIPLA LTD	<u>40MG/2ML (20MG/ML)</u>	<u>A077219 001</u>	Feb 20, 2008
<u>AP</u>			<u>100MG/5ML (20MG/ML)</u>	<u>A077219 002</u>	Feb 20, 2008
<u>AP</u>		EMCURE PHARMS LTD	<u>40MG/2ML (20MG/ML)</u>	<u>A200771 001</u>	Feb 14, 2012
<u>AP</u>			<u>100MG/5ML (20MG/ML)</u>	<u>A200771 002</u>	Feb 14, 2012

PRESCRIPTION DRUG PRODUCT LIST

IRINOTECAN HYDROCHLORIDE

INJECTABLE; INJECTION

IRINOTECAN HYDROCHLORIDE

AP	FRESENIUS KABI USA	<u>40MG/2ML (20MG/ML)</u>	<u>A077776 001</u>	Feb 27, 2008
AP		<u>100MG/5ML (20MG/ML)</u>	<u>A077776 002</u>	Feb 27, 2008
AP	GLAND PHARMA LTD	<u>40MG/2ML (20MG/ML)</u>	<u>A212993 001</u>	Nov 18, 2019
AP		<u>100MG/5ML (20MG/ML)</u>	<u>A212993 002</u>	Nov 18, 2019
AP	HIKMA FARMACEUTICA	<u>40MG/2ML (20MG/ML)</u>	<u>A091032 001</u>	Dec 20, 2010
AP		<u>100MG/5ML (20MG/ML)</u>	<u>A091032 002</u>	Dec 20, 2010
AP	HISUN PHARM HANGZHOU	<u>40MG/2ML (20MG/ML)</u>	<u>A090016 001</u>	Jan 28, 2009
AP		<u>100MG/5ML (20MG/ML)</u>	<u>A090016 002</u>	Jan 28, 2009
AP	HOSPIRA	<u>40MG/2ML (20MG/ML)</u>	<u>A077915 001</u>	Feb 27, 2008
AP		<u>100MG/5ML (20MG/ML)</u>	<u>A077915 002</u>	Feb 27, 2008
AP	!	<u>500MG/2.5ML (20MG/ML)</u>	<u>A078796 001</u>	Feb 27, 2008
AP	INGENUS PHARMS LLC	<u>40MG/2ML (20MG/ML)</u>	<u>A206935 001</u>	May 26, 2017
AP		<u>100MG/5ML (20MG/ML)</u>	<u>A206935 002</u>	May 26, 2017
AP	INTAS PHARMS USA	<u>40MG/2ML (20MG/ML)</u>	<u>A203054 001</u>	Aug 31, 2017
AP		<u>100MG/5ML (20MG/ML)</u>	<u>A203054 002</u>	Aug 31, 2017
AP	JIANGSU HENGRUI MED	<u>40MG/2ML (20MG/ML)</u>	<u>A090675 002</u>	Dec 16, 2011
AP		<u>100MG/5ML (20MG/ML)</u>	<u>A090675 001</u>	Dec 16, 2011
AP	NEOPHARMA	<u>40MG/2ML (20MG/ML)</u>	<u>A078953 001</u>	Apr 15, 2010
AP		<u>100MG/5ML (20MG/ML)</u>	<u>A078953 002</u>	Apr 15, 2010
AP	PLIVA LACHEMA	<u>40MG/2ML (20MG/ML)</u>	<u>A078122 001</u>	Oct 31, 2008
AP		<u>100MG/5ML (20MG/ML)</u>	<u>A078122 002</u>	Oct 31, 2008
AP	QILU	<u>40MG/2ML (20MG/ML)</u>	<u>A203380 001</u>	May 03, 2016
AP		<u>100MG/5ML (20MG/ML)</u>	<u>A203380 002</u>	May 03, 2016
AP		<u>300MG/15ML (20MG/ML)</u>	<u>A203380 003</u>	May 03, 2016
AP	SHILPA MEDICARE LTD	<u>40MG/2ML (20MG/ML)</u>	<u>A208718 001</u>	Dec 28, 2018
AP		<u>100MG/5ML (20MG/ML)</u>	<u>A208718 002</u>	Dec 28, 2018
AP	TEVA PHARMS USA	<u>40MG/2ML (20MG/ML)</u>	<u>A090101 002</u>	Feb 27, 2008
AP		<u>100MG/5ML (20MG/ML)</u>	<u>A090101 003</u>	Feb 27, 2008
AP		<u>500MG/2.5ML (20MG/ML)</u>	<u>A090101 001</u>	Nov 26, 2008
AP	WEST-WARD PHARMS INT	<u>40MG/2ML (20MG/ML)</u>	<u>A078753 001</u>	Dec 24, 2008
AP		<u>100MG/5ML (20MG/ML)</u>	<u>A078753 002</u>	Dec 24, 2008
AP	ZENNOVA	<u>40MG/2ML (20MG/ML)</u>	<u>A090393 002</u>	May 13, 2011
AP		<u>100MG/5ML (20MG/ML)</u>	<u>A090393 003</u>	May 13, 2011

INJECTABLE, LIPOSOMAL; INTRAVENOUS

ONIVYDE

+! IPSEN INC EQ 43MG BASE/10ML (EQ 4.3MG BASE/ML) N207793 001 Oct 22, 2015

IRON DEXTRAN

INJECTABLE; INJECTION

INFED

BP +! ALLERGAN EQ 50MG IRON/ML N017441 001

IRON SUCROSE

INJECTABLE; INTRAVENOUS

VENOFER

+ AM REGENT EQ 50MG BASE/2.5ML (EQ 20MG BASE/ML) N021135 002 Mar 20, 2005

+! EQ 100MG BASE/5ML (EQ 20MG BASE/ML) N021135 001 Nov 06, 2000

+ EQ 200MG BASE/10ML (EQ 20MG BASE/ML) N021135 004 Feb 09, 2007

ISAVUCONAZONIUM SULFATE

CAPSULE; ORAL

CRESEMBA

+! ASTELLAS 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 HLTHCARE 99.9% N017624 001ISOFLURANEAN HALOCARBON PRODS 99.9% A075225 001 Oct 20, 1999AN PIRAMAL CRITICAL 99.9% A074416 001 Sep 30, 1994AN PIRAMAL ENT 99.9% A074502 001 Jun 27, 1995

PRESCRIPTION DRUG PRODUCT LIST

ISONIAZID

INJECTABLE; INJECTION

ISONIAZID

! SANDOZ INC 100MG/ML A040648 001 Jul 05, 2005

SYRUP; ORAL

ISONIAZID

! CMP PHARMA INC 50MG/5ML A088235 001 Nov 10, 1983

TABLET; ORAL

ISONIAZID

<u>AA</u>	ANDA REPOSITORY	<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>	!	<u>100MG</u>	<u>A080936</u>	<u>001</u>	
<u>AA</u>	!	<u>300MG</u>	<u>A080937</u>	<u>002</u>	
<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

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

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>	CIPLA	<u>0.2MG/ML</u>	<u>A210322</u>	<u>001</u>	Jun 12, 2018
<u>AP</u>		<u>0.2MG/ML</u>	<u>A211738</u>	<u>001</u>	Jun 28, 2019
<u>AP</u>	NEXUS PHARMS	<u>0.2MG/ML</u>	<u>A206961</u>	<u>001</u>	Aug 02, 2017

ISUPREL

<u>AP</u>	!	<u>0.2MG/ML</u>	<u>N010515</u>	<u>001</u>	
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ISOSORBIDE DINITRATE

CAPSULE, EXTENDED RELEASE; ORAL

DILATRATE-SR

+! AUXILIUM PHARMS LLC 40MG N019790 001 Sep 02, 1988

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>	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>		<u>30MG</u>	<u>A040591</u>	<u>001</u>	Jan 10, 2007
<u>AB</u>	PAR PHARM	<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>	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	<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

TABLET, EXTENDED RELEASE; ORAL

ISOSORBIDE DINITRATE

! SUN PHARM INDS INC 40MG A040009 001 Dec 30, 1998

ISOSORBIDE MONONITRATE

TABLET; ORAL

ISOSORBIDE MONONITRATE

<u>AB</u>	ACTAVIS ELIZABETH	<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
<u>AB</u>	HIKMA PHARMS	<u>20MG</u>	<u>A075361</u>	<u>001</u>	Oct 05, 2000

PRESCRIPTION DRUG PRODUCT LIST

ISOSORBIDE MONONITRATE

TABLET; ORAL

MONOKET

<u>AB</u>	+	LANNETT CO INC	<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>		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
<u>AB</u>		LANNETT CO INC	<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>		NESHER PHARMS	<u>30MG</u>	<u>A075395</u>	<u>001</u>	Mar 16, 2000
<u>AB</u>			<u>60MG</u>	<u>A075395</u>	<u>002</u>	Mar 16, 2000
<u>AB</u>			<u>120MG</u>	<u>A075395</u>	<u>003</u>	Mar 16, 2000
<u>AB</u>		RICONPHARMA LLC	<u>30MG</u>	<u>A210918</u>	<u>001</u>	Nov 05, 2018
<u>AB</u>			<u>60MG</u>	<u>A210918</u>	<u>002</u>	Nov 05, 2018
<u>AB</u>			<u>120MG</u>	<u>A210918</u>	<u>003</u>	Nov 05, 2018
<u>AB</u>		TORRENT PHARMS	<u>30MG</u>	<u>A200270</u>	<u>001</u>	Jun 03, 2011
<u>AB</u>			<u>60MG</u>	<u>A200495</u>	<u>001</u>	Jun 03, 2011
<u>AB</u>			<u>120MG</u>	<u>A200495</u>	<u>002</u>	Jun 03, 2011
<u>AB</u>		VINTAGE PHARMS	<u>30MG</u>	<u>A090598</u>	<u>001</u>	Aug 11, 2010
<u>AB</u>			<u>60MG</u>	<u>A090598</u>	<u>002</u>	Aug 11, 2010
<u>AB</u>			<u>120MG</u>	<u>A090598</u>	<u>003</u>	Aug 11, 2010

ISOSULFAN BLUE

INJECTABLE; INJECTION

ISOSULFAN BLUE

<u>AP</u>		AUROBINDO PHARMA LTD	<u>1%</u>	<u>A206831</u>	<u>001</u>	Feb 02, 2016
<u>AP</u>		!	<u>1%</u>	<u>A090874</u>	<u>001</u>	Jul 20, 2010

ISOTRETINOIN

CAPSULE; ORAL

AMNESTEEM

<u>AB</u>		MYLAN PHARMS INC	<u>10MG</u>	<u>A075945</u>	<u>001</u>	Nov 08, 2002
<u>AB</u>			<u>20MG</u>	<u>A075945</u>	<u>002</u>	Nov 08, 2002
<u>AB</u>			<u>40MG</u>	<u>A075945</u>	<u>003</u>	Nov 08, 2002

CLARAVIS

<u>AB</u>		TEVA PHARMS USA	<u>10MG</u>	<u>A076356</u>	<u>001</u>	Apr 11, 2003
<u>AB</u>			<u>20MG</u>	<u>A076135</u>	<u>002</u>	Apr 11, 2003
<u>AB</u>			<u>30MG</u>	<u>A076135</u>	<u>003</u>	May 11, 2006
<u>AB</u>		!	<u>40MG</u>	<u>A076135</u>	<u>001</u>	Apr 11, 2003

ISOTRETINOIN

<u>AB</u>		AMNEAL PHARMS NY	<u>10MG</u>	<u>A207792</u>	<u>001</u>	Sep 29, 2017
<u>AB</u>			<u>20MG</u>	<u>A207792</u>	<u>002</u>	Sep 29, 2017
<u>AB</u>			<u>30MG</u>	<u>A207792</u>	<u>003</u>	Sep 29, 2017
<u>AB</u>			<u>40MG</u>	<u>A207792</u>	<u>004</u>	Sep 29, 2017

MYORISAN

<u>AB</u>		DOUGLAS PHARMS	<u>10MG</u>	<u>A076485</u>	<u>001</u>	Jan 19, 2012
<u>AB</u>			<u>20MG</u>	<u>A076485</u>	<u>002</u>	Jan 19, 2012
<u>AB</u>			<u>30MG</u>	<u>A076485</u>	<u>004</u>	Aug 25, 2015
<u>AB</u>			<u>40MG</u>	<u>A076485</u>	<u>003</u>	Jan 19, 2012

ZENATANE

<u>AB</u>		DR REDDYS LABS LTD	<u>10MG</u>	<u>A202099</u>	<u>001</u>	Mar 25, 2013
<u>AB</u>			<u>20MG</u>	<u>A202099</u>	<u>002</u>	Mar 25, 2013
<u>AB</u>			<u>30MG</u>	<u>A202099</u>	<u>004</u>	Feb 23, 2015
<u>AB</u>			<u>40MG</u>	<u>A202099</u>	<u>003</u>	Mar 25, 2013

ABSORICA

BX	+	SUN PHARM INDS INC	10MG	N021951	001	May 25, 2012
BX	+		20MG	N021951	002	May 25, 2012
BX	+		30MG	N021951	003	May 25, 2012
BX	+	!	40MG	N021951	004	May 25, 2012
	+		25MG	N021951	005	Aug 15, 2014
	+		35MG	N021951	006	Aug 15, 2014

ABSORICA LD

	+	SUN PHARM	8MG	N211913	001	Nov 05, 2019
	+		16MG	N211913	002	Nov 05, 2019
	+		20MG	N211913	003	Nov 05, 2019
	+		24MG	N211913	004	Nov 05, 2019
	+		28MG	N211913	005	Nov 05, 2019
	+	!	32MG	N211913	006	Nov 05, 2019

PRESCRIPTION DRUG PRODUCT LISTISRADIPINE

CAPSULE; ORAL

ISRADIPINE

AB	ELITE LABS INC	2.5MG	A077169 001	Apr 24, 2006
AB		5MG	A077169 002	Apr 24, 2006
AB	WATSON LABS TEVA	2.5MG	A077317 001	Jan 05, 2006
AB	!	5MG	A077317 002	Jan 05, 2006

ISTRADÉFYLLINE

TABLET; ORAL

NOURIANZ

+	KYOWA KIRIN	20MG	N022075 001	Aug 27, 2019
+	!	40MG	N022075 002	Aug 27, 2019

ITRACONAZOLE

CAPSULE; ORAL

ITRACONAZOLE

AB	ACCORD HLTHCARE	100MG	A205991 001	May 26, 2016
AB	ALEMBIC PHARMS LTD	100MG	A206741 001	Dec 13, 2016
AB	ALKEM LABS LTD	100MG	A208591 001	Jun 12, 2017
AB	AMNEAL PHARMS	100MG	A205080 001	Sep 26, 2016
AB	JUBILANT GENERICS	100MG	A203445 001	Feb 23, 2017
AB	MYLAN PHARMS INC	100MG	A200463 001	Jul 20, 2012
AB	PAR PHARM INC	100MG	A205724 001	Dec 13, 2016
AB	PII	100MG	A206410 001	Jul 02, 2019
AB	SANDOZ	100MG	A076104 001	May 28, 2004
AB	TORRENT	100MG	A209460 001	Aug 24, 2018
AB	ZYDUS PHARMS	100MG	A204672 001	Sep 19, 2017

SPORANOX

AB	+	JANSSEN PHARMS	100MG	N020083 001	Sep 11, 1992
		TOLSURA			
	+	MAYNE PHARMA INTL	65MG	N208901 001	Dec 11, 2018

SOLUTION; ORAL

ITRACONAZOLE

AA	AMNEAL PHARMS	10MG/ML	A205573 001	Oct 30, 2015
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SPORANOX

AA	+	JANSSEN PHARMS	10MG/ML	N020657 001	Feb 21, 1997
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TABLET; ORAL

ONMEL

+	SEBELA IRELAND LTD	200MG	N022484 001	Apr 29, 2010
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IVABRADINE

SOLUTION; ORAL

CORLANOR

+	AMGEN INC	5MG/5ML (1MG/ML)	N209964 001	Apr 22, 2019
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IVABRADINE HYDROCHLORIDE

TABLET; ORAL

CORLANOR

+	AMGEN INC	EQ 5MG BASE	N206143 001	Apr 15, 2015
+	!	EQ 7.5MG BASE	N206143 002	Apr 15, 2015

IVACAFTOR

GRANULE; ORAL

KALYDECO

+	VERTEX PHARMS INC	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	75MG	N203188 002	May 20, 2019
+	!	150MG	N203188 001	Jan 31, 2012

IVACAFTOR; IVACAFTOR, TEZACAFTOR

TABLET, TABLET; ORAL

SYMDEKO (COPACKAGED)

+	VERTEX PHARMS INC	75MG, N/A; 75MG, 50MG	N210491 002	Jun 21, 2019
+	!	150MG, N/A; 150MG, 100MG	N210491 001	Feb 12, 2018

IVACAFTOR; LUMACAFTOR

GRANULE; ORAL

ORKAMBI

+	VERTEX PHARMS INC	125MG/PACKET; 100MG/PACKET	N211358 001	Aug 07, 2018
+	!	188MG/PACKET; 150MG/PACKET	N211358 002	Aug 07, 2018

PRESCRIPTION DRUG PRODUCT LIST

3-246 (of 453)

IVACAFTOR; LUMACAFTOR

TABLET; ORAL

ORKAMBI

+ VERTEX PHARMS INC 125MG;100MG
 +! 125MG;200MG

N206038 002 Sep 28, 2016
 N206038 001 Jul 02, 2015

IVERMECTIN

CREAM; TOPICAL

IVERMECTIN**AB** TEVA PHARMS USA **1%****A210019 001** Sep 13, 2019SOOLANTRA**AB** +! GALDERMA LABS LP **1%****N206255 001** Dec 19, 2014

LOTION; TOPICAL

SKLICE

+! ARBOR PHARMS LLC 0.5%

N202736 001 Feb 07, 2012

TABLET; ORAL

IVERMECTIN**AB** EDENBRIDGE PHARMS **3MG****A204154 001** Oct 24, 2014STROMEKTOL**AB** +! MERCK SHARP DOHME **3MG****N050742 002** Oct 08, 1998IVOSIDENIB

TABLET; ORAL

TIBSOVO

+! AGIOS PHARMS INC 250MG

N211192 001 Jul 20, 2018

IXABEPILONE

INJECTABLE; INTRAVENOUS

IXEMPRA KIT

+! R-PHARM US LLC 15MG/VIAL
 +! 45MG/VIAL

N022065 001 Oct 16, 2007
 N022065 002 Oct 16, 2007

IXAZOMIB CITRATE

CAPSULE; ORAL

NINLARO

+ MILLENNIUM PHARMS EQ 2.3MG BASE
 + EQ 3MG BASE
 +! EQ 4MG BASE

N208462 001 Nov 20, 2015
 N208462 002 Nov 20, 2015
 N208462 003 Nov 20, 2015

KETAMINE HYDROCHLORIDE

INJECTABLE; INJECTION

KETALAR**AP** +! PAR STERILE PRODUCTS **EQ 10MG BASE/ML****N016812 001****AP** +! **EQ 50MG BASE/ML****N016812 002****AP** +! **EQ 100MG BASE/ML****N016812 003**KETAMINE HYDROCHLORIDE**AP** HOSPIRA **EQ 50MG BASE/ML****A074549 001** Jun 27, 1996**AP** **EQ 100MG BASE/ML****A074549 002** Jun 27, 1996**AP** MYLAN INSTITUTIONAL **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** WEST-WARD PHARMS **EQ 50MG BASE/ML****A074524 001** Mar 22, 1996

INT

AP **EQ 100MG BASE/ML****A074524 002** Mar 22, 1996KETOCONAZOLE

AEROSOL, FOAM; TOPICAL

EXTINA**AT** +! MYLAN **2%****N021738 001** Jun 12, 2007KETOCONAZOLE**AT** PERRIGO ISRAEL **2%****A091550 001** Aug 25, 2011

CREAM; TOPICAL

KETOCONAZOLE**AB** FOUGERA PHARMS **2%****A076294 001** Apr 28, 2004**AB** ! TEVA **2%****A075581 001** Apr 25, 2000KETOZOLE**AB** TARO **2%****A075638 001** Dec 18, 2002

GEL; TOPICAL

XOLEGEL

+! ALMIRALL 2%

N021946 001 Jul 28, 2006

SHAMPOO; TOPICAL

KETOCONAZOLE**AB** PERRIGO ISRAEL **2%****A076419 001** Jan 07, 2004**AB** TOLMAR **2%****A076942 001** Apr 11, 2005

PRESCRIPTION DRUG PRODUCT LIST

KETOCONAZOLE

SHAMPOO; TOPICAL

NIZORAL

AB +! JANSSEN PHARMS **2%** **N019927 001** Aug 31, 1990
 TABLET; ORAL

KETOCONAZOLE

AB MYLAN **200MG** **A075597 001** Dec 23, 1999
AB STRIDES PHARMA **200MG** **A210457 001** Jun 18, 2018
AB ! TARO **200MG** **A075319 001** Jun 15, 1999

KETOPROFEN

CAPSULE; ORAL

KETOPROFEN

AB HERITAGE PHARMS INC **50MG** **A074014 002** Jan 29, 1993
AB **75MG** **A074014 003** Jan 29, 1993
AB TEVA **50MG** **A073516 001** Dec 22, 1992
AB ! **75MG** **A073517 001** Dec 22, 1992
 HERITAGE PHARMS INC 25MG A074014 001 Jan 29, 1993

CAPSULE, EXTENDED RELEASE; ORAL

KETOPROFEN

! MYLAN 200MG A075679 001 Feb 20, 2002

KETOROLAC TROMETHAMINE

INJECTABLE; INJECTION

KETOROLAC TROMETHAMINE

AP AMPHASTAR PHARM **15MG/ML** **A076209 001** Jul 21, 2004
AP **30MG/ML** **A076209 002** Jul 21, 2004
AP BAXTER HLTHCARE **15MG/ML** **A209900 002** Jul 25, 2018
 CORP
AP **30MG/ML** **A209900 001** Sep 15, 2017
AP FRESENIUS KABI USA **15MG/ML** **A075784 001** Jan 11, 2002
AP **15MG/ML** **A203242 001** Oct 07, 2015
AP **30MG/ML** **A075784 002** Jan 11, 2002
AP **30MG/ML** **A203242 002** Oct 07, 2015
AP GLAND PHARMA LTD **15MG/ML** **A204216 001** Nov 01, 2016
AP **30MG/ML** **A204216 002** Nov 01, 2016
AP ! HOSPIRA **15MG/ML** **A074802 001** Jun 05, 1997
AP **15MG/ML** **A074993 001** Jan 27, 1999
AP ! **30MG/ML** **A074802 002** Jun 05, 1997
AP **30MG/ML** **A074993 002** Jan 27, 1999
AP SAGENT PHARMS INC **15MG/ML** **A091065 001** Nov 27, 2013
AP **30MG/ML** **A091065 002** Nov 27, 2013
AP SANDOZ INC **30MG/ML** **A076271 002** Oct 06, 2004
AP WEST-WARD PHARMS **15MG/ML** **A075772 001** Jul 21, 2004
 INT
AP **30MG/ML** **A075772 002** Jul 21, 2004
AP WOCKHARDT **15MG/ML** **A077942 001** Mar 27, 2007
AP **30MG/ML** **A077942 002** Mar 27, 2007

SOLUTION/DROPS; OPHTHALMIC

ACULAR

AT +! ALLERGAN **0.5%** **N019700 001** Nov 09, 1992

ACULAR LS

AT +! ALLERGAN **0.4%** **N021528 001** May 30, 2003

KETOROLAC TROMETHAMINE

AT AKORN **0.4%** **A078399 001** Nov 05, 2009
AT **0.5%** **A078434 001** Nov 05, 2009
AT APOTEX INC **0.4%** **A077308 001** Nov 05, 2009
AT **0.5%** **A076109 001** Nov 05, 2009
AT MICRO LABS LTD **0.5%** **A203410 001** Apr 05, 2019
 INDIA
AT SANDOZ INC **0.5%** **A076583 001** Nov 05, 2009
AT SUN PHARM **0.5%** **A090017 001** Nov 05, 2009

ACUVAIL

+! ALLERGAN 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 CYCLE PHARMS LTD **10MG** **A210616 001** Aug 16, 2018
AB MYLAN **10MG** **A074761 001** May 16, 1997
AB PLIVA **10MG** **A075284 001** Jun 23, 1999
AB ! TEVA **10MG** **A074754 001** May 16, 1997

PRESCRIPTION DRUG PRODUCT LIST

KETOROLAC TROMETHAMINE; PHENYLEPHRINE HYDROCHLORIDE

SOLUTION;IRRIGATION

KETOROLAC TROMETHAMINE AND PHENYLEPHRINE HYDROCHLORIDE

AT	LUPIN LTD	<u>EQ 0.3% BASE;EQ 1% BASE</u>	<u>A210183 001</u>	Jul 01, 2019
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OMIDRIA

AT	+! OMEROS	<u>EQ 0.3% BASE;EQ 1% BASE</u>	<u>N205388 001</u>	May 30, 2014
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L-GLUTAMINE

FOR SOLUTION;ORAL

ENDARI

	+ EMMAUS MEDCL	5GM/PACKET	N208587 001	Jul 07, 2017
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LABETALOL HYDROCHLORIDE

INJECTABLE;INJECTION

LABETALOL HYDROCHLORIDE

AP	AKORN INC	<u>5MG/ML</u>	<u>A075431 001</u>	Nov 29, 1999
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AP	GLAND PHARMA LTD	<u>5MG/ML</u>	<u>A090699 001</u>	Apr 03, 2012
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AP	! HOSPIRA	<u>5MG/ML</u>	<u>A075239 001</u>	Nov 29, 1999
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AP	!	<u>5MG/ML</u>	<u>A075240 001</u>	Nov 29, 1999
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AP	WEST-WARD PHARMS INT	<u>5MG/ML</u>	<u>A075303 001</u>	May 28, 1999
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TABLET;ORAL

LABETALOL HYDROCHLORIDE

AB	Athem	<u>100MG</u>	<u>A207863 001</u>	Feb 04, 2019
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AB		<u>200MG</u>	<u>A207863 002</u>	Feb 04, 2019
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AB		<u>300MG</u>	<u>A207863 003</u>	Feb 04, 2019
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AB	CADILA PHARMS LTD	<u>100MG</u>	<u>A211325 001</u>	May 13, 2019
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AB		<u>200MG</u>	<u>A211325 002</u>	May 13, 2019
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AB		<u>300MG</u>	<u>A211325 003</u>	May 13, 2019
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AB	HERITAGE PHARMA	<u>100MG</u>	<u>A074787 001</u>	Aug 03, 1998
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AB		<u>200MG</u>	<u>A074787 002</u>	Aug 03, 1998
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AB	INNOGENIX	<u>100MG</u>	<u>A075215 001</u>	Jul 29, 1999
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AB		<u>200MG</u>	<u>A075215 002</u>	Jul 29, 1999
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AB		<u>300MG</u>	<u>A075215 003</u>	Jul 29, 1999
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AB	PAR FORM	<u>100MG</u>	<u>A200908 001</u>	Jul 10, 2012
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AB		<u>200MG</u>	<u>A200908 002</u>	Jul 10, 2012
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AB		<u>300MG</u>	<u>A200908 003</u>	Jul 10, 2012
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AB	SANDOZ	<u>100MG</u>	<u>A075113 001</u>	Aug 04, 1998
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AB	!	<u>200MG</u>	<u>A075113 002</u>	Aug 04, 1998
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AB		<u>300MG</u>	<u>A075113 003</u>	Aug 04, 1998
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AB	TWI PHARMS	<u>100MG</u>	<u>A209603 001</u>	Jun 20, 2018
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AB		<u>200MG</u>	<u>A209603 002</u>	Jun 20, 2018
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AB		<u>300MG</u>	<u>A209603 003</u>	Jun 20, 2018
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AB	WATSON LABS	<u>100MG</u>	<u>A075133 001</u>	Aug 03, 1998
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AB		<u>200MG</u>	<u>A075133 002</u>	Aug 03, 1998
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AB		<u>300MG</u>	<u>A075133 003</u>	Aug 03, 1998
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AB	ZYDUS PHARMS	<u>100MG</u>	<u>A207743 001</u>	Sep 19, 2017
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AB		<u>200MG</u>	<u>A207743 002</u>	Sep 19, 2017
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AB		<u>300MG</u>	<u>A207743 003</u>	Sep 19, 2017
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TRANDATE

AB	+ CNTY LINE PHARMS	<u>100MG</u>	<u>N018716 001</u>	May 24, 1985
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AB	+	<u>200MG</u>	<u>N018716 002</u>	Aug 01, 1984
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AB	+	<u>300MG</u>	<u>N018716 003</u>	Aug 01, 1984
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LACOSAMIDE

SOLUTION;INTRAVENOUS

VIMPAT

	+! UCB INC	200MG/20ML (10MG/ML)	N022254 001	Oct 28, 2008
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SOLUTION;ORAL

VIMPAT

	+! UCB INC	10MG/ML	N022255 001	Apr 20, 2010
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TABLET;ORAL

VIMPAT

	+ UCB INC	50MG	N022253 001	Oct 28, 2008
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	+	100MG	N022253 002	Oct 28, 2008
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	+	150MG	N022253 003	Oct 28, 2008
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	+!	200MG	N022253 004	Oct 28, 2008
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LACTITOL

FOR SOLUTION;ORAL

PIZENSY

	+! BRAINTREE LABS	10GM	N211281 001	Feb 12, 2020
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PRESCRIPTION DRUG PRODUCT LIST

LACTULOSE

FOR SOLUTION;ORAL

LACTULOSE

!	CUMBERLAND PHARMS	10GM/PACKET	A074712	001	Dec 10, 1997
!		20GM/PACKET	A074712	002	Dec 10, 1997

SOLUTION;ORAL

CONSTILAC

<u>AA</u>	ALRA	<u>10GM/15ML</u>	<u>A071054</u>	<u>001</u>	Jul 26, 1988
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LACTULOSE

<u>AA</u>	FRESENIUS KABI	<u>10GM/15ML</u>	<u>A090503</u>	<u>001</u>	Jan 25, 2012
<u>AA</u>	! HI TECH PHARMA	<u>10GM/15ML</u>	<u>A074076</u>	<u>001</u>	Jul 03, 1995
<u>AA</u>	HIKMA	<u>10GM/15ML</u>	<u>A073591</u>	<u>001</u>	May 29, 1992
<u>AA</u>	LANNETT CO INC	<u>10GM/15ML</u>	<u>A075993</u>	<u>001</u>	Jul 26, 2001
<u>AA</u>	LIFEPHARMA	<u>10GM/15ML</u>	<u>A209517</u>	<u>001</u>	Nov 23, 2018
<u>AA</u>	PHARM ASSOC	<u>10GM/15ML</u>	<u>A074623</u>	<u>001</u>	Jul 30, 1996
<u>AA</u>	VISTAPHARM	<u>10GM/15ML</u>	<u>A074138</u>	<u>001</u>	Sep 30, 1992
<u>AA</u>	WOCKHARDT BIO AG	<u>10GM/15ML</u>	<u>A074602</u>	<u>001</u>	Nov 14, 1996
<u>AA</u>	XTTRIUM LABS INC	<u>10GM/15ML</u>	<u>A075911</u>	<u>001</u>	Feb 21, 2002

SOLUTION;ORAL, RECTAL

CHOLAC

<u>AA</u>	ALRA	<u>10GM/15ML</u>	<u>A071331</u>	<u>001</u>	Jul 26, 1988
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ENULOSE

<u>AA</u>	! ACTAVIS MID ATLANTIC	<u>10GM/15ML</u>	<u>A071548</u>	<u>001</u>	Aug 15, 1988
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GENERLAC

<u>AA</u>	WOCKHARDT BIO AG	<u>10GM/15ML</u>	<u>A074603</u>	<u>001</u>	Oct 31, 1996
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LACTULOSE

<u>AA</u>	FRESENIUS KABI	<u>10GM/15ML</u>	<u>A090502</u>	<u>001</u>	Jan 25, 2012
<u>AA</u>	HI TECH PHARMA	<u>10GM/15ML</u>	<u>A074077</u>	<u>001</u>	Jul 03, 1995

LAMIVUDINE

SOLUTION;ORAL

EPIVIR

<u>AA</u>	+! VIIV HLTHCARE	<u>10MG/ML</u>	<u>N020596</u>	<u>001</u>	Nov 17, 1995
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LAMIVUDINE

<u>AA</u>	AUROBINDO PHARMA LTD	<u>10MG/ML</u>	<u>A077695</u>	<u>001</u>	Nov 21, 2016
<u>AA</u>	LANNETT CO INC	<u>10MG/ML</u>	<u>A203564</u>	<u>001</u>	Oct 31, 2014

EPIVIR-HBV

+	GLAXOSMITHKLINE	5MG/ML	N021004	001	Dec 08, 1998
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TABLET;ORAL

EPIVIR

<u>AB</u>	+ VIIV HLTHCARE	<u>150MG</u>	<u>N020564</u>	<u>001</u>	Nov 17, 1995
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<u>AB</u>	+!	<u>300MG</u>	<u>N020564</u>	<u>003</u>	Jun 24, 2002
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EPIVIR-HBV

<u>AB</u>	+! GLAXOSMITHKLINE	<u>100MG</u>	<u>N021003</u>	<u>001</u>	Dec 08, 1998
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LAMIVUDINE

<u>AB</u>	ANNORA	<u>100MG</u>	<u>A211306</u>	<u>001</u>	Mar 21, 2019
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<u>AB</u>	APOTEX	<u>100MG</u>	<u>A202941</u>	<u>001</u>	Jan 02, 2014
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<u>AB</u>		<u>150MG</u>	<u>A091606</u>	<u>001</u>	Dec 02, 2011
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<u>AB</u>		<u>300MG</u>	<u>A091606</u>	<u>002</u>	Dec 02, 2011
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<u>AB</u>	ARISE	<u>150MG</u>	<u>A206974</u>	<u>001</u>	Nov 21, 2016
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<u>AB</u>		<u>300MG</u>	<u>A206974</u>	<u>002</u>	Nov 21, 2016
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<u>AB</u>	AUROBINDO PHARMA LTD	<u>150MG</u>	<u>A077464</u>	<u>001</u>	Nov 21, 2016
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<u>AB</u>		<u>150MG</u>	<u>A202032</u>	<u>001</u>	Nov 17, 2011
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<u>AB</u>		<u>300MG</u>	<u>A077464</u>	<u>002</u>	Nov 21, 2016
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<u>AB</u>		<u>300MG</u>	<u>A202032</u>	<u>002</u>	Nov 17, 2011
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<u>AB</u>	CIPLA	<u>150MG</u>	<u>A077221</u>	<u>001</u>	Mar 03, 2017
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<u>AB</u>		<u>300MG</u>	<u>A077221</u>	<u>002</u>	Mar 03, 2017
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<u>AB</u>	ECI PHARMS LLC	<u>150MG</u>	<u>A203586</u>	<u>001</u>	Nov 21, 2016
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<u>AB</u>	HETERO LABS LTD V	<u>100MG</u>	<u>A203260</u>	<u>001</u>	Jan 02, 2014
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<u>AB</u>		<u>150MG</u>	<u>A203277</u>	<u>001</u>	Jan 06, 2014
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<u>AB</u>		<u>300MG</u>	<u>A203277</u>	<u>002</u>	Jan 06, 2014
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<u>AB</u>	LUPIN LTD	<u>150MG</u>	<u>A205217</u>	<u>001</u>	Dec 18, 2014
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<u>AB</u>		<u>300MG</u>	<u>A205217</u>	<u>002</u>	Dec 18, 2014
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<u>AB</u>	MACLEODS PHARMS LTD	<u>150MG</u>	<u>A090198</u>	<u>001</u>	May 01, 2019
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<u>AB</u>		<u>300MG</u>	<u>A090198</u>	<u>002</u>	May 01, 2019
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<u>AB</u>	MYLAN LABS LTD	<u>150MG</u>	<u>A078545</u>	<u>001</u>	Mar 05, 2019
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<u>AB</u>		<u>300MG</u>	<u>A078545</u>	<u>002</u>	Mar 05, 2019
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<u>AB</u>	STRIDES PHARMA	<u>150MG</u>	<u>A090457</u>	<u>001</u>	Apr 19, 2018
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<u>AB</u>		<u>300MG</u>	<u>A090457</u>	<u>002</u>	Apr 19, 2018
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PRESCRIPTION DRUG PRODUCT LISTLAMIVUDINE; TENOFOVIR DISOPROXIL FUMARATE

TABLET; ORAL

CIMDUO

+! MYLAN LABS LTD 300MG;300MG N022141 001 Feb 28, 2018

TEMIXYS

CELLTRION 300MG;300MG N211284 001 Nov 16, 2018

LAMIVUDINE; ZIDOVUDINE

TABLET; ORAL

COMBIVIR**AB +! VIIV HLTHCARE 150MG;300MG N020857 001 Sep 26, 1997****LAMIVUDINE AND ZIDOVUDINE****AB ANDA REPOSITORY 150MG;300MG A206375 001 Apr 10, 2018****AB AUROBINDO PHARMA LTD 150MG;300MG A077558 001 May 05, 2017****AB 150MG;300MG A202418 001 May 15, 2012****AB CIPLA 150MG;300MG A077411 001 Sep 07, 2018****AB HETERO LABS LTD III 150MG;300MG A079124 001 Sep 17, 2015****AB HETERO LABS LTD V 150MG;300MG A203259 001 Feb 03, 2014****AB LUPIN LTD 150MG;300MG A090246 001 May 15, 2012****AB MACLEODS PHARMS LTD 150MG;300MG A090679 001 Aug 29, 2018****AB STRIDES PHARMA 150MG;300MG A079128 001 May 13, 2015**LAMOTRIGINE

TABLET; ORAL

LAMICTAL**AB +! GLAXOSMITHKLINE LLC 25MG N020241 005 Dec 27, 1994****AB + 100MG N020241 001 Dec 27, 1994****AB + 150MG N020241 002 Dec 27, 1994****AB + 200MG N020241 003 Dec 27, 1994****LAMOTRIGINE****AB ALEMBIC PHARMS LTD 25MG A090607 001 Jan 13, 2011****AB 100MG A090607 002 Jan 13, 2011****AB 150MG A090607 003 Jan 13, 2011****AB 200MG A090607 004 Jan 13, 2011****AB ALKEM LABS LTD 25MG A200694 001 Jun 14, 2013****AB 100MG A200694 002 Jun 14, 2013****AB 150MG A200694 003 Jun 14, 2013****AB 200MG A200694 004 Jun 14, 2013****AB AUROBINDO PHARMA 25MG A078956 001 Jan 27, 2009****AB 100MG A078956 002 Jan 27, 2009****AB 150MG A078956 003 Jan 27, 2009****AB 200MG A078956 004 Jan 27, 2009****AB CIPLA 25MG A077783 001 Nov 01, 2010****AB 100MG A077783 002 Nov 01, 2010****AB 150MG A077783 003 Nov 01, 2010****AB 200MG A077783 004 Nov 01, 2010****AB DR REDDYS LABS LTD 25MG A076708 001 Jan 27, 2009****AB 100MG A076708 002 Jan 27, 2009****AB 150MG A076708 003 Jan 27, 2009****AB 200MG A076708 004 Jan 27, 2009****AB GLENMARK GENERICS 25MG A090169 001 May 12, 2012****AB 100MG A090169 002 May 12, 2012****AB 150MG A090169 003 May 12, 2012****AB 200MG A090169 004 May 12, 2012****AB JUBILANT CADISTA 25MG A079132 001 Jan 27, 2009****AB 100MG A079132 002 Jan 27, 2009****AB 150MG A079132 003 Jan 27, 2009****AB 200MG A079132 004 Jan 27, 2009****AB LUPIN LTD 25MG A078691 001 Jun 01, 2010****AB 100MG A078691 002 Jun 01, 2010****AB 150MG A078691 003 Jun 01, 2010****AB 200MG A078691 004 Jun 01, 2010****AB MYLAN 25MG A077420 001 Jan 27, 2009****AB 100MG A077420 002 Jan 27, 2009****AB 150MG A077420 003 Jan 27, 2009****AB 200MG A077420 004 Jan 27, 2009****AB RUBICON 25MG A078625 001 Jan 27, 2009****AB 100MG A078625 002 Jan 27, 2009****AB 150MG A078625 003 Jan 27, 2009****AB 200MG A078625 004 Jan 27, 2009****AB TARO PHARM INDS 25MG A078525 001 Jan 27, 2009****AB 100MG A078525 002 Jan 27, 2009****AB 150MG A078525 003 Jan 27, 2009**

PRESCRIPTION DRUG PRODUCT LIST

LAMOTRIGINE

TABLET;ORAL

LAMOTRIGINE

<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, CHEWABLE;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 PHARMS LTD	<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>	JUBILANT GENERICS	<u>5MG</u>	<u>A200220 001</u>	Feb 28, 2011
<u>AB</u>		<u>25MG</u>	<u>A200220 002</u>	Feb 28, 2011
<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, 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>A203733 001</u>	Nov 13, 2013
<u>AB</u>		<u>300MG</u>	<u>A200672 005</u>	Oct 17, 2013
<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
<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>	ANCHEN PHARMS	<u>25MG</u>	<u>A201374 001</u>	Dec 26, 2012
<u>AB</u>		<u>50MG</u>	<u>A201374 002</u>	Dec 26, 2012
<u>AB</u>		<u>100MG</u>	<u>A201374 003</u>	Dec 26, 2012
<u>AB</u>		<u>200MG</u>	<u>A201374 004</u>	Dec 26, 2012
<u>AB</u>		<u>250MG</u>	<u>A201374 005</u>	Dec 26, 2012
<u>AB</u>		<u>300MG</u>	<u>A201374 006</u>	Dec 26, 2012
<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

PRESCRIPTION DRUG PRODUCT LIST

LAMOTRIGINE

TABLET, EXTENDED RELEASE;ORAL

LAMOTRIGINE

<u>AB</u>		<u>300MG</u>	<u>A202383</u>	<u>005</u>	Jun 19, 2013
<u>AB</u>	PAR PHARM	<u>25MG</u>	<u>A201791</u>	<u>001</u>	Jan 18, 2013
<u>AB</u>		<u>50MG</u>	<u>A201791</u>	<u>002</u>	Jan 18, 2013
<u>AB</u>		<u>100MG</u>	<u>A201791</u>	<u>003</u>	Jan 18, 2013
<u>AB</u>		<u>200MG</u>	<u>A201791</u>	<u>004</u>	Jan 18, 2013
<u>AB</u>		<u>250MG</u>	<u>A201791</u>	<u>005</u>	Jan 18, 2013
<u>AB</u>		<u>300MG</u>	<u>A201791</u>	<u>006</u>	Jan 18, 2013
<u>AB</u>	RUBICON	<u>100MG</u>	<u>A202887</u>	<u>003</u>	Jun 17, 2013
<u>AB</u>		<u>200MG</u>	<u>A202887</u>	<u>004</u>	Jun 17, 2013
<u>AB</u>	TORRENT	<u>25MG</u>	<u>A203370</u>	<u>001</u>	Dec 23, 2013
<u>AB</u>		<u>50MG</u>	<u>A203370</u>	<u>002</u>	Dec 23, 2013
<u>AB</u>		<u>100MG</u>	<u>A203370</u>	<u>003</u>	Dec 23, 2013
<u>AB</u>		<u>200MG</u>	<u>A203370</u>	<u>004</u>	Dec 23, 2013
<u>AB</u>	WOCKHARDT LTD	<u>25MG</u>	<u>A202498</u>	<u>001</u>	Jan 04, 2013
<u>AB</u>		<u>50MG</u>	<u>A202498</u>	<u>002</u>	Jan 04, 2013
<u>AB</u>		<u>100MG</u>	<u>A202498</u>	<u>003</u>	Jan 04, 2013
<u>AB</u>		<u>200MG</u>	<u>A202498</u>	<u>004</u>	Jan 04, 2013
<u>AB</u>		<u>300MG</u>	<u>A202498</u>	<u>005</u>	Jan 04, 2013

TABLET, ORALLY DISINTEGRATING;ORAL

LAMICTAL ODT

<u>AB</u>	+	GLAXOSMITHKLINE LLC	<u>25MG</u>	<u>N022251</u>	<u>001</u>	May 08, 2009
<u>AB</u>	+	!	<u>50MG</u>	<u>N022251</u>	<u>002</u>	May 08, 2009
<u>AB</u>	+		<u>100MG</u>	<u>N022251</u>	<u>003</u>	May 08, 2009
<u>AB</u>	+		<u>200MG</u>	<u>N022251</u>	<u>004</u>	May 08, 2009

LAMOTRIGINE

<u>AB</u>	IMPAX LABS INC	<u>25MG</u>	<u>A200828</u>	<u>001</u>	Jul 15, 2013
<u>AB</u>		<u>50MG</u>	<u>A200828</u>	<u>002</u>	Jul 15, 2013
<u>AB</u>		<u>100MG</u>	<u>A200828</u>	<u>003</u>	Jul 15, 2013
<u>AB</u>		<u>200MG</u>	<u>A200828</u>	<u>004</u>	Jul 15, 2013
<u>AB</u>	PAR PHARM	<u>25MG</u>	<u>A204158</u>	<u>001</u>	Oct 27, 2015
<u>AB</u>		<u>50MG</u>	<u>A204158</u>	<u>002</u>	Oct 27, 2015
<u>AB</u>		<u>100MG</u>	<u>A204158</u>	<u>003</u>	Oct 27, 2015
<u>AB</u>		<u>200MG</u>	<u>A204158</u>	<u>004</u>	Oct 27, 2015
<u>AB</u>	SCIEGEN PHARMS INC	<u>25MG</u>	<u>A206382</u>	<u>001</u>	Jun 17, 2016
<u>AB</u>		<u>50MG</u>	<u>A206382</u>	<u>002</u>	Jun 17, 2016
<u>AB</u>		<u>100MG</u>	<u>A206382</u>	<u>003</u>	Jun 17, 2016
<u>AB</u>		<u>200MG</u>	<u>A206382</u>	<u>004</u>	Jun 17, 2016

LANREOTIDE ACETATE

SOLUTION;SUBCUTANEOUS

SOMATULINE DEPOT

+	!	IPSEN PHARMA	EQ 60MG BASE/0.2ML (EQ 60MG BASE/0.2ML)	N022074	001	Aug 30, 2007
+	!		EQ 90MG BASE/0.3ML (EQ 90MG BASE/0.3ML)	N022074	002	Aug 30, 2007
+	!		EQ 120MG BASE/0.5ML (EQ 120MG BASE/0.5ML)	N022074	003	Aug 30, 2007

LANSOPRAZOLE

CAPSULE, DELAYED REL PELLETS;ORAL

LANSOPRAZOLE

<u>AB</u>	ALKEM LABS LTD	<u>15MG</u>	<u>A207394</u>	<u>001</u>	Jan 18, 2019
<u>AB</u>		<u>30MG</u>	<u>A207394</u>	<u>002</u>	Jan 18, 2019
<u>AB</u>	DR REDDYS LABS LTD	<u>15MG</u>	<u>A091269</u>	<u>001</u>	Oct 15, 2010
<u>AB</u>		<u>30MG</u>	<u>A091269</u>	<u>002</u>	Oct 15, 2010
<u>AB</u>	INVENTIA	<u>15MG</u>	<u>A205868</u>	<u>001</u>	Aug 30, 2017
<u>AB</u>		<u>30MG</u>	<u>A205868</u>	<u>002</u>	Aug 30, 2017
<u>AB</u>	LANNETT CO INC	<u>15MG</u>	<u>A207156</u>	<u>001</u>	Sep 28, 2017
<u>AB</u>		<u>30MG</u>	<u>A207156</u>	<u>002</u>	Sep 28, 2017
<u>AB</u>	MYLAN PHARMS INC	<u>15MG</u>	<u>A090763</u>	<u>001</u>	Nov 10, 2009
<u>AB</u>		<u>30MG</u>	<u>A090763</u>	<u>002</u>	Nov 10, 2009
<u>AB</u>	NATCO PHARMA LTD	<u>15MG</u>	<u>A201921</u>	<u>001</u>	Dec 18, 2012
<u>AB</u>		<u>30MG</u>	<u>A201921</u>	<u>002</u>	Dec 18, 2012
<u>AB</u>	SANDOZ	<u>15MG</u>	<u>A090331</u>	<u>001</u>	Apr 23, 2010
<u>AB</u>		<u>30MG</u>	<u>A090331</u>	<u>002</u>	Apr 23, 2010
<u>AB</u>	SUN PHARM	<u>15MG</u>	<u>A202637</u>	<u>001</u>	Sep 13, 2013
<u>AB</u>		<u>30MG</u>	<u>A091509</u>	<u>001</u>	Sep 13, 2013
<u>AB</u>	TEVA PHARMS	<u>15MG</u>	<u>A077255</u>	<u>001</u>	Nov 10, 2009
<u>AB</u>		<u>30MG</u>	<u>A077255</u>	<u>002</u>	Nov 10, 2009
<u>AB</u>	WOCKHARDT USA	<u>15MG</u>	<u>A202176</u>	<u>001</u>	Sep 14, 2012
<u>AB</u>		<u>30MG</u>	<u>A202176</u>	<u>002</u>	Sep 14, 2012
<u>AB</u>	XIROMED	<u>15MG</u>	<u>A203203</u>	<u>001</u>	Jul 25, 2016
<u>AB</u>		<u>30MG</u>	<u>A203203</u>	<u>002</u>	Jul 25, 2016

PRESCRIPTION DRUG PRODUCT LIST

LANSOPRAZOLE

CAPSULE, DELAYED REL PELLETS;ORAL

LANSOPRAZOLE

AB	ZYDUS HLTHCARE	15MG	A202366 001	Aug 19, 2013
AB		30MG	A202366 002	Aug 19, 2013

PREVACID

AB	+ TAKEDA PHARMS USA	15MG	N020406 001	May 10, 1995
AB	+!	30MG	N020406 002	May 10, 1995

TABLET, ORALLY DISINTEGRATING, DELAYED RELEASE;ORAL

LANSOPRAZOLE

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

PREVACID

AB	+ TAKEDA PHARMS USA	15MG	N021428 001	Aug 30, 2002
AB	+!	30MG	N021428 002	Aug 30, 2002

LANTHANUM CARBONATE

POWDER;ORAL

FOSRENOL

	+ SHIRE DEV LLC	EQ 750MG BASE	N204734 001	Sep 24, 2014
	+!	EQ 1GM BASE	N204734 002	Sep 24, 2014

TABLET, CHEWABLE;ORAL

FOSRENOL

AB	+ SHIRE LLC	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	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

TYKERB

	+! NOVARTIS	EQ 250MG BASE	N022059 001	Mar 13, 2007
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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
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LASMITAN SUCCINATE

TABLET;ORAL

REYVOW

	+ ELI LILLY AND CO	50MG	N211280 001	Jan 31, 2020
	+!	100MG	N211280 002	Jan 31, 2020

LATANOPROST

EMULSION;OPHTHALMIC

XELPROS

	+! SUN PHARMA GLOBAL	0.005%	N206185 001	Sep 12, 2018
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SOLUTION/DROPS;OPHTHALMIC

LATANOPROST

AT	AKORN	0.005%	A090887 001	Jul 19, 2011
AT	AMRING PHARMS	0.005%	A200925 001	Mar 22, 2011
AT	AUROBINDO PHARMA LTD	0.005%	A206519 001	Sep 03, 2019
AT	BAUSCH AND LOMB	0.005%	A201006 001	Mar 22, 2011
AT	DR REDDYS LABS LTD	0.005%	A202077 001	Feb 11, 2013
AT	FDC LTD	0.005%	A202442 001	Apr 22, 2016
AT	SANDOZ INC	0.005%	A091449 001	Mar 22, 2011
AT	SOMERSET	0.005%	A201786 001	Mar 22, 2011

XALATAN

AT	+! PHARMACIA AND UPJOHN	0.005%	N020597 001	Jun 05, 1996
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PRESCRIPTION DRUG PRODUCT LISTLATANOPROST; NETARSUDIL DIMESYLATE

SOLUTION/DROPS;OPHTHALMIC

ROCKLATAN

+! AERIE PHARMS 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

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

+! NABRIVA EQ 150MG BASE/15ML (EQ 10MG BASE/ML) N211673 001 Aug 19, 2019

TABLET;ORAL

XENLETA

+! NABRIVA 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** 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** BARR **10MG****A077083 001** Sep 13, 2005**AB** **20MG****A077083 002** Sep 13, 2005**AB** HERITAGE PHARMS INC **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** TEVA PHARMS **10MG****A077084 001** Sep 13, 2005**AB** **20MG****A077084 002** Sep 13, 2005**AB** ZYDUS **10MG****A212308 001** Apr 24, 2019**AB** **20MG****A212308 002** Apr 24, 2019

ARAVA

+! SANOFI AVENTIS US 100MG N020905 003 Sep 10, 1998

LENALIDOMIDE

CAPSULE;ORAL

REVLIMID

+ CELGENE 2.5MG N021880 005 Dec 21, 2011

+ 5MG N021880 001 Dec 27, 2005

+ 10MG N021880 002 Dec 27, 2005

+ 15MG N021880 003 Jun 29, 2006

+ 20MG N021880 006 Jun 05, 2013

+! 25MG N021880 004 Jun 29, 2006

LENVATINIB MESYLATE

CAPSULE;ORAL

LENVIMA

+ EISAI INC EQ 4MG BASE N206947 001 Feb 13, 2015

+! EQ 10MG BASE N206947 002 Feb 13, 2015

LETERMOVIR

SOLUTION;INTRAVENOUS

PREVYMIS

+! MERCK SHARP DOHME 240MG/12ML (20MG/ML) N209940 001 Nov 08, 2017

+! 480MG/24ML (20MG/ML) N209940 002 Nov 08, 2017

PRESCRIPTION DRUG PRODUCT LIST

LETERMOVIR

TABLET; ORAL

PREVYMIS

+	MERCK SHARP DOHME	240MG	N209939	001	Nov 08, 2017
+	!	480MG	N209939	002	Nov 08, 2017

LETROZOLE

TABLET; ORAL

FEMARA

AB	+	NOVARTIS PHARMS	2.5MG	N020726	001	Jul 25, 1997
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LETROZOLE

AB		ACCORD HLTHCARE	2.5MG	A090934	001	Jun 03, 2011
AB		APOTEX INC	2.5MG	A091303	001	Apr 19, 2012
AB		BEIJING YILING	2.5MG	A205869	001	Nov 14, 2018
AB		DR REDDYS LABS LTD	2.5MG	A091191	001	Jun 03, 2011
AB		EUGIA PHARMA	2.5MG	A211717	001	Jan 11, 2019
AB		HIKMA	2.5MG	A090838	001	Jun 03, 2011
AB		HIKMA PHARMS	2.5MG	A203796	001	Jun 03, 2016
AB		INDICUS PHARMA	2.5MG	A201804	001	Jun 03, 2011
AB		JIANGSU HENGRUI MED	2.5MG	A202716	001	May 16, 2013
AB		NATCO PHARMA LTD	2.5MG	A200161	001	Jun 03, 2011
AB		TEVA PHARMS	2.5MG	A090289	001	Jun 03, 2011
AB		VINTAGE PHARMS LLC	2.5MG	A090789	001	Jun 03, 2011

LETROZOLE; RIBOCICLIB SUCCINATE

TABLET, TABLET; ORAL

KISQALI FEMARA CO-PACK (COPACKAGED)

+	NOVARTIS	2.5MG,N/A;N/A,EQ 200MG BASE	N209935	001	May 04, 2017
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LEUCOVORIN CALCIUM

INJECTABLE; INJECTION

LEUCOVORIN CALCIUM

AP		FRESENIUS KABI USA	EQ 10MG BASE/ML	A207226	001	Jul 27, 2018
AP		TEVA PHARMS USA	EQ 100MG BASE/VIAL	A081277	001	Sep 28, 1993
AP			EQ 350MG BASE/VIAL	A040174	001	Jun 12, 1997
AP	!	WEST-WARD PHARMS INT	EQ 50MG BASE/VIAL	A089384	001	Sep 14, 1987
AP	!		EQ 100MG BASE/VIAL	A089717	001	Mar 28, 1988

LEUCOVORIN CALCIUM PRESERVATIVE FREE

AP		FRESENIUS KABI USA	EQ 200MG BASE/VIAL	A040258	001	Feb 26, 1999
AP	!		EQ 500MG BASE/VIAL	A040286	001	Feb 26, 1999
AP		MYLAN LABS LTD	EQ 100MG BASE/VIAL	A203800	001	May 19, 2017
AP			EQ 200MG BASE/VIAL	A203800	002	May 19, 2017
AP			EQ 350MG BASE/VIAL	A203800	003	May 19, 2017
AP		SAGENT PHARMS	EQ 50MG BASE/VIAL	A200753	001	Sep 06, 2012
AP			EQ 100MG BASE/VIAL	A200753	002	Sep 06, 2012
AP			EQ 200MG BASE/VIAL	A200753	003	Sep 06, 2012
AP			EQ 350MG BASE/VIAL	A200855	001	Sep 06, 2012
AP		SAGENT PHARMS INC	EQ 500MG BASE/VIAL	A209110	001	Oct 26, 2017
AP	!	WEST-WARD PHARMS INT	EQ 10MG BASE/ML	A040347	001	Apr 25, 2000
AP	!		EQ 200MG BASE/VIAL	A040056	001	May 23, 1995
AP	!		EQ 350MG BASE/VIAL	A040335	001	Apr 20, 2000

LEUCOVORIN CALCIUM

FRESENIUS KABI USA EQ 10MG BASE/ML

A207241 001 Mar 14, 2018

TABLET; ORAL

LEUCOVORIN CALCIUM

AB		BARR	EQ 5MG BASE	A071198	001	Sep 24, 1987
AB			EQ 25MG BASE	A071199	001	Sep 24, 1987
AB		HIKMA	EQ 5MG BASE	A072733	001	Feb 22, 1993
AB	!		EQ 25MG BASE	A072736	001	Feb 22, 1993
			EQ 10MG BASE	A072734	001	Feb 22, 1993
			EQ 15MG BASE	A072735	001	Feb 22, 1993

LEUPROLIDE ACETATE

INJECTABLE; INJECTION

LEUPROLIDE ACETATE

AP	!	SANDOZ	1MG/0.2ML	A074728	001	Aug 04, 1998
AP		SUN PHARM	1MG/0.2ML	A078885	001	Mar 09, 2009
AP		TEVA PHARMS USA	1MG/0.2ML	A075471	001	Oct 25, 2000

LUPRON DEPOT

+	ABBVIE ENDOCRINE INC	3.75MG	N020011	002	Oct 26, 1995
+		7.5MG/VIAL	N019732	001	Jan 26, 1989
+		11.25MG/VIAL	N020708	001	Mar 07, 1997

PRESCRIPTION DRUG PRODUCT LISTLEUPROLIDE ACETATE

INJECTABLE; INJECTION

LUPRON DEPOT

+		22.5MG/VIAL	N020517	001	Dec 22, 1995
+	!	30MG/VIAL	N020517	002	May 30, 1997
+	!	45MG/VIAL	N020517	003	Jun 17, 2011

LUPRON DEPOT-PED

+	!	ABBVIE ENDOCRINE INC	7.5MG/VIAL	N020263	002	Apr 16, 1993
+	!		11.25MG/VIAL	N020263	005	Jan 21, 1994
+	!		11.25MG/VIAL	N020263	007	Aug 15, 2011
+	!		15MG/VIAL	N020263	006	Jan 21, 1994
+	!		30MG/VIAL	N020263	008	Aug 15, 2011

INJECTABLE; SUBCUTANEOUS

ELIGARD

+	!	TOLMAR THERAP	7.5MG/VIAL	N021343	001	Jan 23, 2002
+	!		22.5MG/VIAL	N021379	001	Jul 24, 2002
+	!		30MG/VIAL	N021488	001	Feb 13, 2003
+	!		45MG/VIAL	N021731	001	Dec 14, 2004

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

<u>AN</u>		AUROBINDO PHARMA LTD	<u>EQ 0.25% BASE</u>	<u>A207628</u>	<u>001</u>	Jan 31, 2017
<u>AN</u>			<u>EQ 0.0103% BASE</u>	<u>A207625</u>	<u>001</u>	Dec 30, 2016
<u>AN</u>			<u>EQ 0.021% BASE</u>	<u>A207625</u>	<u>002</u>	Dec 30, 2016
<u>AN</u>			<u>EQ 0.042% BASE</u>	<u>A207625</u>	<u>003</u>	Dec 30, 2016
<u>AN</u>		CIPLA	<u>EQ 0.021% BASE</u>	<u>A078171</u>	<u>002</u>	Dec 13, 2013
<u>AN</u>			<u>EQ 0.042% BASE</u>	<u>A078171</u>	<u>003</u>	Dec 13, 2013
<u>AN</u>			<u>EQ 0.0103% BASE</u>	<u>A078171</u>	<u>001</u>	Dec 13, 2013
<u>AN</u>		IMPAX LABS INC	<u>EQ 0.0103% BASE</u>	<u>A077756</u>	<u>003</u>	Apr 09, 2008
<u>AN</u>			<u>EQ 0.021% BASE</u>	<u>A077756</u>	<u>001</u>	Apr 09, 2008
<u>AN</u>			<u>EQ 0.042% BASE</u>	<u>A077756</u>	<u>002</u>	Apr 09, 2008
<u>AN</u>		MYLAN SPECIALITY LP	<u>EQ 0.25% BASE</u>	<u>A078309</u>	<u>001</u>	Mar 20, 2009
<u>AN</u>		RITEDOSE CORP	<u>EQ 0.0103% BASE</u>	<u>A203653</u>	<u>001</u>	Mar 22, 2016
<u>AN</u>			<u>EQ 0.021% BASE</u>	<u>A203653</u>	<u>002</u>	Mar 22, 2016
<u>AN</u>			<u>EQ 0.042% BASE</u>	<u>A203653</u>	<u>003</u>	Mar 22, 2016
<u>AN</u>		SUN PHARM	<u>EQ 0.0103% BASE</u>	<u>A207820</u>	<u>001</u>	Nov 05, 2018
<u>AN</u>			<u>EQ 0.021% BASE</u>	<u>A207820</u>	<u>002</u>	Nov 05, 2018
<u>AN</u>			<u>EQ 0.042% BASE</u>	<u>A207820</u>	<u>003</u>	Nov 05, 2018
<u>AN</u>		TEVA PARENTERAL	<u>EQ 0.25% BASE</u>	<u>A200875</u>	<u>001</u>	Sep 11, 2014
<u>AN</u>		TEVA PHARMS USA	<u>EQ 0.0103% BASE</u>	<u>A090297</u>	<u>001</u>	Apr 26, 2013
<u>AN</u>			<u>EQ 0.021% BASE</u>	<u>A090297</u>	<u>002</u>	Apr 26, 2013
<u>AN</u>			<u>EQ 0.042% BASE</u>	<u>A090297</u>	<u>003</u>	Apr 26, 2013

XOPENEX

<u>AN</u>	+	OAK PHARMS INC	<u>EQ 0.0103% BASE</u>	<u>N020837</u>	<u>003</u>	Jan 30, 2002
<u>AN</u>	+		<u>EQ 0.021% BASE</u>	<u>N020837</u>	<u>001</u>	Mar 25, 1999
<u>AN</u>	+		<u>EQ 0.042% BASE</u>	<u>N020837</u>	<u>002</u>	Mar 25, 1999
<u>AN</u>	+		<u>EQ 0.25% BASE</u>	<u>N020837</u>	<u>004</u>	Jul 18, 2003

LEVALBUTEROL TARTRATE

AEROSOL, METERED; INHALATION

XOPENEX HFA

+	!	SUNOVION	EQ 0.045MG BASE/INH	N021730	001	Mar 11, 2005
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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

PRESCRIPTION DRUG PRODUCT LIST

LEVETIRACETAM

INJECTABLE; INTRAVENOUS

KEPPRA

<u>AP</u>	<u>+!</u> UCB INC	<u>500MG/5ML (100MG/ML)</u>	<u>N021872 001</u>	Jul 31, 2006
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LEVETIRACETAM

<u>AP</u>	AUROBINDO PHARMA LTD	<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>A090813 001</u>	May 26, 2010
<u>AP</u>		<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>	JUBILANT GENERICS	<u>500MG/5ML (100MG/ML)</u>	<u>A206838 001</u>	Jun 02, 2016
<u>AP</u>	MICRO LABS	<u>500MG/5ML (100MG/ML)</u>	<u>A211954 001</u>	Aug 09, 2019
<u>AP</u>	MYLAN LABS LTD	<u>500MG/5ML (100MG/ML)</u>	<u>A203308 001</u>	Sep 16, 2016
<u>AP</u>	SAGENT PHARMS	<u>500MG/5ML (100MG/ML)</u>	<u>A091627 001</u>	Jun 26, 2013
<u>AP</u>	SUN PHARM INDS LTD	<u>500MG/5ML (100MG/ML)</u>	<u>A090754 001</u>	Jun 16, 2010
<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>	AUROBINDO PHARMA LTD	<u>500MG/100ML (5MG/ML)</u>	<u>A207160 001</u>	Jan 04, 2017
<u>AP</u>		<u>1000MG/100ML (10MG/ML)</u>	<u>A207160 002</u>	Jan 04, 2017
<u>AP</u>		<u>1500MG/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>1000MG/100ML (10MG/ML)</u>	<u>A206880 002</u>	Oct 25, 2017
<u>AP</u>		<u>1500MG/100ML (15MG/ML)</u>	<u>A206880 003</u>	Oct 25, 2017
<u>AP</u>	<u>+!</u> HQ SPECIALITY PHARMA	<u>500MG/100ML (5MG/ML)</u>	<u>N202543 001</u>	Nov 09, 2011
<u>AP</u>	<u>+!</u>	<u>1000MG/100ML (10MG/ML)</u>	<u>N202543 002</u>	Nov 09, 2011
<u>AP</u>	<u>+!</u>	<u>1500MG/100ML (15MG/ML)</u>	<u>N202543 003</u>	Nov 09, 2011

SOLUTION; ORAL

KEPPRA

<u>AA</u>	<u>+!</u> UCB INC	<u>100MG/ML</u>	<u>N021505 001</u>	Jul 15, 2003
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LEVETIRACETAM

<u>AA</u>	ACTAVIS MID ATLANTIC	<u>100MG/ML</u>	<u>A078976 001</u>	Jan 15, 2009
<u>AA</u>	AMNEAL PHARMS	<u>100MG/ML</u>	<u>A090992 001</u>	Oct 27, 2009
<u>AA</u>	ANDA REPOSITORY	<u>100MG/ML</u>	<u>A079107 001</u>	Jan 15, 2009
<u>AA</u>	AUROBINDO PHARMA LTD	<u>100MG/ML</u>	<u>A079063 001</u>	Jan 15, 2009
<u>AA</u>	BRECKENRIDGE	<u>100MG/ML</u>	<u>A079120 001</u>	Jan 16, 2009
<u>AA</u>	HETERO LABS LTD III	<u>100MG/ML</u>	<u>A203052 001</u>	Feb 28, 2013
<u>AA</u>	HI-TECH PHARMACAL	<u>100MG/ML</u>	<u>A090601 001</u>	Feb 28, 2012
<u>AA</u>	LANNETT CO INC	<u>100MG/ML</u>	<u>A090079 001</u>	Apr 11, 2012
<u>AA</u>		<u>100MG/ML</u>	<u>A090263 001</u>	Apr 03, 2009
<u>AA</u>	LUPIN LTD	<u>100MG/ML</u>	<u>A090893 001</u>	Oct 17, 2011
<u>AA</u>	ORIT LABS LLC	<u>100MG/ML</u>	<u>A203067 001</u>	May 09, 2013
<u>AA</u>	PHARM ASSOC	<u>100MG/ML</u>	<u>A201157 001</u>	Jun 04, 2015
<u>AA</u>	TARO	<u>100MG/ML</u>	<u>A078774 001</u>	Feb 10, 2009
<u>AA</u>	TRIS PHARMA INC	<u>100MG/ML</u>	<u>A090461 001</u>	Sep 30, 2010
<u>AA</u>	WOCKHARDT BIO AG	<u>100MG/ML</u>	<u>A090028 001</u>	Mar 03, 2010

TABLET; ORAL

KEPPRA

<u>AB</u>	<u>+</u> UCB INC	<u>250MG</u>	<u>N021035 001</u>	Nov 30, 1999
<u>AB</u>	<u>+</u>	<u>500MG</u>	<u>N021035 002</u>	Nov 30, 1999
<u>AB</u>	<u>+</u>	<u>750MG</u>	<u>N021035 003</u>	Nov 30, 1999
<u>AB</u>	<u>+!</u>	<u>1GM</u>	<u>N021035 004</u>	Jan 06, 2006

LEVETIRACETAM

<u>AB</u>	ACCORD HLTHCARE	<u>250MG</u>	<u>A090843 001</u>	Feb 14, 2011
<u>AB</u>		<u>500MG</u>	<u>A090843 002</u>	Feb 14, 2011
<u>AB</u>		<u>750MG</u>	<u>A090843 003</u>	Feb 14, 2011
<u>AB</u>		<u>1GM</u>	<u>A090843 004</u>	Feb 14, 2011
<u>AB</u>	ACI HEALTHCARE LTD	<u>250MG</u>	<u>A078042 001</u>	Jan 15, 2009
<u>AB</u>		<u>500MG</u>	<u>A078042 002</u>	Jan 15, 2009
<u>AB</u>		<u>750MG</u>	<u>A078042 003</u>	Jan 15, 2009
<u>AB</u>		<u>1GM</u>	<u>A078042 004</u>	Jan 15, 2009
<u>AB</u>	ACIC PHARMS	<u>250MG</u>	<u>A090767 001</u>	Jul 28, 2010
<u>AB</u>		<u>500MG</u>	<u>A090767 002</u>	Jul 28, 2010
<u>AB</u>		<u>750MG</u>	<u>A090767 003</u>	Jul 28, 2010
<u>AB</u>		<u>1GM</u>	<u>A090767 004</u>	Jul 28, 2010
<u>AB</u>	APOTEX INC	<u>250MG</u>	<u>A078869 001</u>	Mar 13, 2009
<u>AB</u>		<u>500MG</u>	<u>A078869 002</u>	Mar 13, 2009
<u>AB</u>		<u>750MG</u>	<u>A078869 003</u>	Mar 13, 2009

PRESCRIPTION DRUG PRODUCT LIST

LEVETIRACETAM

TABLET; ORAL

LEVETIRACETAM

<u>AB</u>		<u>1GM</u>	<u>A078869</u>	<u>004</u>	Mar 13, 2009
<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>	BRECKENRIDGE PHARM	<u>250MG</u>	<u>A090511</u>	<u>001</u>	Aug 18, 2011
<u>AB</u>		<u>500MG</u>	<u>A090511</u>	<u>002</u>	Aug 18, 2011
<u>AB</u>		<u>750MG</u>	<u>A090511</u>	<u>003</u>	Aug 18, 2011
<u>AB</u>		<u>1GM</u>	<u>A090511</u>	<u>004</u>	Aug 18, 2011
<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>	INVAGEN PHARMS	<u>250MG</u>	<u>A078234</u>	<u>001</u>	Jan 15, 2009
<u>AB</u>		<u>500MG</u>	<u>A078234</u>	<u>002</u>	Jan 15, 2009
<u>AB</u>		<u>750MG</u>	<u>A078234</u>	<u>003</u>	Jan 15, 2009
<u>AB</u>	LOTUS PHARM CO LTD	<u>250MG</u>	<u>A090906</u>	<u>002</u>	Oct 31, 2016
<u>AB</u>		<u>500MG</u>	<u>A090906</u>	<u>001</u>	Nov 05, 2010
<u>AB</u>		<u>750MG</u>	<u>A090906</u>	<u>003</u>	Oct 31, 2016
<u>AB</u>		<u>1GM</u>	<u>A090906</u>	<u>004</u>	Oct 31, 2016
<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>	MYLAN	<u>250MG</u>	<u>A076919</u>	<u>001</u>	Nov 04, 2008
<u>AB</u>		<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>	ORCHID HLTHCARE	<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
<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>	SECAN PHARMS	<u>500MG</u>	<u>A205102</u>	<u>004</u>	Dec 16, 2015
<u>AB</u>		<u>1GM</u>	<u>A205102</u>	<u>003</u>	Dec 16, 2015
<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>	TEVA PHARMS	<u>250MG</u>	<u>A078101</u>	<u>001</u>	Jan 15, 2009
<u>AB</u>		<u>500MG</u>	<u>A078101</u>	<u>002</u>	Jan 15, 2009
<u>AB</u>		<u>750MG</u>	<u>A078101</u>	<u>003</u>	Jan 15, 2009
<u>AB</u>		<u>1GM</u>	<u>A078101</u>	<u>004</u>	Jan 15, 2009
<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>	VINTAGE PHARMS	<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>	WOCKHARDT	<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

PRESCRIPTION DRUG PRODUCT LIST

LEVETIRACETAM

TABLET;ORAL

LEVETIRACETAM

AB	ZYDUS PHARMS USA INC	250MG	A078918 001	Apr 29, 2009
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AB		1GM	A078918 002	Apr 29, 2009
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TABLET, EXTENDED RELEASE;ORAL

KEPPRA XR

AB	+ UCB INC	500MG	N022285 001	Sep 12, 2008
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AB	+	750MG	N022285 002	Feb 12, 2009
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LEVETIRACETAM

AB	ACTAVIS ELIZABETH	500MG	A091557 001	Sep 12, 2011
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AB		750MG	A091557 002	Sep 12, 2011
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AB	ACTAVIS LABS FL INC	500MG	A091093 001	Sep 12, 2011
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AB		750MG	A091093 002	Sep 12, 2011
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AB	ANCHEN PHARMS	500MG	A091360 001	Oct 04, 2011
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AB		750MG	A091360 002	Oct 04, 2011
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AB	ANDA REPOSITORY	500MG	A204511 001	Feb 23, 2016
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AB		750MG	A204511 002	Feb 23, 2016
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AB	APOTEX INC	500MG	A091261 001	Sep 12, 2011
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AB		750MG	A091261 002	Sep 12, 2011
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AB	DEXCEL PHARMA	500MG	A202167 001	Sep 04, 2015
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AB		750MG	A202167 002	Sep 04, 2015
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AB	ECI PHARMS LLC	500MG	A204754 001	Aug 26, 2016
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AB		750MG	A204754 002	Aug 26, 2016
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AB	HISUN PHARM HANGZHOU	500MG	A207175 001	Sep 28, 2017
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AB		750MG	A207175 002	Sep 28, 2017
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AB	LOTUS PHARM CO LTD	500MG	A202095 002	Jun 06, 2016
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AB		750MG	A202095 001	Jun 06, 2016
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AB	LUPIN LTD	500MG	A091399 001	Sep 12, 2011
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AB		750MG	A091399 002	Sep 12, 2011
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AB	PHARMADAX INC	500MG	A201464 001	May 25, 2012
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AB		750MG	A201464 002	May 25, 2012
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AB	PRINSTON INC	500MG	A202533 001	Jul 20, 2012
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AB		500MG	A203468 001	May 21, 2015
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AB		750MG	A202533 002	Jul 20, 2012
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AB		750MG	A203468 002	May 21, 2015
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AB	ROUSES POINT PHARMS	500MG	A202524 001	Aug 27, 2012
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AB		750MG	A202524 002	Aug 27, 2012
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AB	SUN PHARM	500MG	A203059 001	Sep 09, 2013
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AB		750MG	A203059 002	Sep 09, 2013
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AB	TEVA PHARMS	500MG	A091430 001	Sep 12, 2011
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AB		750MG	A091430 002	Sep 12, 2011
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AB	TORRENT PHARMS LTD	500MG	A091338 001	May 29, 2012
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AB		750MG	A091338 002	May 29, 2012
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	ELEPSIA XR			
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	+ SPARC	1GM	N204417 001	Dec 20, 2018
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	+	1.5GM	N204417 002	Dec 20, 2018
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	LEVETIRACETAM			
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	APOTEX	1GM	A202958 001	Feb 25, 2015
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TABLET, FOR SUSPENSION;ORAL

	SPRITAM			
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	+ APRECIA PHARMS	250MG	N207958 001	Jul 31, 2015
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	+	500MG	N207958 002	Jul 31, 2015
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	+	750MG	N207958 003	Jul 31, 2015
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	+	1GM	N207958 004	Jul 31, 2015
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	LEVOBUNOLOL HYDROCHLORIDE			
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SOLUTION/DROPS;OPHTHALMIC

	<u>AKBETA</u>			
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AT	AKORN	0.25%	A074779 001	Oct 29, 1996
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AT		0.5%	A074780 001	Oct 29, 1996
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	<u>BETAGAN</u>			
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AT	+	0.25%	N019814 001	Jun 28, 1989
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AT	+	0.5%	N019219 002	Dec 19, 1985
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	<u>LEVOBUNOLOL HYDROCHLORIDE</u>			
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AT	BAUSCH AND LOMB	0.5%	A074326 001	Mar 04, 1994
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PRESCRIPTION DRUG PRODUCT LIST

LEVOCARNITINE

INJECTABLE; INJECTION

CARNITOR

AP	+	LEADIANT BIOSCI INC	200MG/ML	N020182	001	Dec 16, 1992
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LEVOCARNITINE

AP		AM REGENT	200MG/ML	A075861	001	Jun 22, 2001
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AP		WEST-WARD PHARMS	200MG/ML	A075567	001	Mar 29, 2001
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INT

SOLUTION; ORAL

CARNITOR

AA	+	LEADIANT BIOSCI INC	1GM/10ML	N019257	001	Apr 10, 1986
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CARNITOR SF

AA	+	LEADIANT BIOSCI INC	1GM/10ML	N019257	002	Mar 28, 2007
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LEVOCARNITINE

AA		HI TECH PHARMA	1GM/10ML	A077399	001	Oct 25, 2007
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AA		LYNE	1GM/10ML	A076851	001	Aug 10, 2004
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AA		NOVITIUM PHARMA	1GM/10ML	A211676	001	Aug 14, 2019
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LEVOCARNITINE SF

AA		NOVITIUM PHARMA	1GM/10ML	A211676	002	Aug 14, 2019
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TABLET; ORAL

CARNITOR

AB	+	LEADIANT BIOSCI INC	330MG	N018948	001	Dec 27, 1985
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LEVOCARNITINE

AB		RISING	330MG	A076858	001	Sep 20, 2004
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LEVOCETIRIZINE DIHYDROCHLORIDE

SOLUTION; ORAL

LEVOCETIRIZINE DIHYDROCHLORIDE

AA	!	L PERRIGO CO	2.5MG/5ML	A091263	001	Nov 07, 2011
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AA		LANNETT CO INC	2.5MG/5ML	A204599	001	May 15, 2017
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AA		TARO PHARM INDS LTD	2.5MG/5ML	A202673	001	Jul 26, 2013
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LEVOCETIRIZINE HYDROCHLORIDE

AA		HETERO LABS LTD III	2.5MG/5ML	A210914	001	Apr 01, 2019
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XYZAL

AA	+	SANOFI AVENTIS US	2.5MG/5ML	N022157	001	Jan 28, 2008
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TABLET; ORAL

LEVOCETIRIZINE DIHYDROCHLORIDE

AB		DR REDDYS LABS LTD	5MG	A090392	001	Feb 24, 2011
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AB		GLENMARK GENERICS	5MG	A090385	001	Feb 24, 2011
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AB		HETERO LABS LTD III	5MG	A091264	001	Jun 29, 2012
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AB		MACLEODS PHARMS LTD	5MG	A205564	001	Jan 11, 2016
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AB		MICRO LABS LTD	5MG	A202046	001	Sep 17, 2013
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INDIA

AB		NEOPHARMA	5MG	A204323	001	Dec 20, 2016
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AB		SCIEGEN PHARMS INC	5MG	A203646	001	Sep 09, 2014
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AB		SUN PHARM	5MG	A090362	001	Jan 31, 2013
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AB		SUN PHARM INDS LTD	5MG	A201653	001	Jun 26, 2015
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AB		SYNTHON PHARMS	5MG	A090229	001	Nov 26, 2010
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AB		TEVA PHARMS	5MG	A090199	001	Aug 22, 2011
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XYZAL

AB	+	SANOFI AVENTIS US	5MG	N022064	001	May 25, 2007
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LEVODOPA

POWDER; INHALATION

INBRIJA

+	!	ACORDA	42MG	N209184	001	Dec 21, 2018
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LEVOFLOXACIN

INJECTABLE; INJECTION

LEVOFLOXACIN

AP		AKORN	EQ 500MG/20ML (EQ 25MG/ML)	A091644	001	Jun 20, 2011
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AP			EQ 750MG/30ML (EQ 25MG/ML)	A091644	002	Jun 20, 2011
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AP	!	AUROBINDO PHARMA LTD	EQ 500MG/20ML (EQ 25MG/ML)	A202328	001	Jan 24, 2013
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AP	!		EQ 750MG/30ML (EQ 25MG/ML)	A202328	002	Jan 24, 2013
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AP		BAXTER HLTHCARE CORP	EQ 500MG/20ML (EQ 25MG/ML)	A091436	001	Jun 05, 2013
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LEVOFLOXACIN IN DEXTROSE 5% IN PLASTIC CONTAINER

AP		AUROBINDO PHARMA LTD	EQ 250MG/50ML (EQ 5MG/ML)	A206919	001	Feb 10, 2016
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AP			EQ 500MG/100ML (EQ 5MG/ML)	A206919	002	Feb 10, 2016
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AP			EQ 750MG/150ML (EQ 5MG/ML)	A206919	003	Feb 10, 2016
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AP		BAXTER HLTHCARE CORP	EQ 250MG/50ML (EQ 5MG/ML)	A091397	001	Aug 08, 2013
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AP			EQ 500MG/100ML (EQ 5MG/ML)	A091397	002	Aug 08, 2013
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PRESCRIPTION DRUG PRODUCT LIST

LEVOFLOXACIN

INJECTABLE; INJECTION

LEVOFLOXACIN IN DEXTROSE 5% IN PLASTIC CONTAINER

<u>AP</u>		<u>EQ 750MG/150ML (EQ 5MG/ML)</u>	<u>A091397 003</u>	Aug 08, 2013
<u>AP</u>	FRESENIUS KABI USA	<u>EQ 250MG/50ML (EQ 5MG/ML)</u>	<u>A200674 001</u>	Jun 19, 2013
<u>AP</u>		<u>EQ 500MG/100ML (EQ 5MG/ML)</u>	<u>A200674 002</u>	Jun 19, 2013
<u>AP</u>		<u>EQ 750MG/150ML (EQ 5MG/ML)</u>	<u>A200674 003</u>	Jun 19, 2013
<u>AP</u>	HIKMA FARMACEUTICA	<u>EQ 250MG/50ML (EQ 5MG/ML)</u>	<u>A091375 001</u>	Sep 16, 2011
<u>AP</u>		<u>EQ 500MG/100ML (EQ 5MG/ML)</u>	<u>A091375 002</u>	Sep 16, 2011
<u>AP</u>		<u>EQ 750MG/150ML (EQ 5MG/ML)</u>	<u>A091375 003</u>	Sep 16, 2011
<u>AP</u>	HOSPIRA INC	<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>	!	<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>	!	HI TECH PHARMA	<u>250MG/10ML</u>	<u>A091678 001</u>	Jun 20, 2011
<u>AA</u>		LANNETT CO INC	<u>250MG/10ML</u>	<u>A205222 001</u>	May 25, 2018

SOLUTION/DROPS; OPHTHALMIC

LEVOFLOXACIN

<u>AT</u>		AKORN	<u>0.5%</u>	<u>A090268 001</u>	Dec 20, 2010
<u>AT</u>		MYLAN LABS LTD	<u>0.5%</u>	<u>A204899 001</u>	Dec 08, 2017
<u>AT</u>	!	RISING	<u>0.5%</u>	<u>A077700 001</u>	Dec 20, 2010
<u>AT</u>		WATSON LABS TEVA	<u>0.5%</u>	<u>A076826 001</u>	Feb 10, 2011
		MICRO LABS LTD	<u>1.5%</u>	<u>A205600 001</u>	Feb 27, 2019
		INDIA			

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>		CELLTRION	<u>250MG</u>	<u>A090367 001</u>	Jun 20, 2011
<u>AB</u>			<u>500MG</u>	<u>A090367 002</u>	Jun 20, 2011
<u>AB</u>			<u>750MG</u>	<u>A090367 003</u>	Jun 20, 2011
<u>AB</u>		CIPLA LTD	<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 GENERICS	<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>		JUBILANT GENERICS	<u>250MG</u>	<u>A203613 001</u>	Jun 19, 2015
<u>AB</u>			<u>500MG</u>	<u>A203613 002</u>	Jun 19, 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>		ORCHID HLTHCARE	<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>		SANDOZ	<u>250MG</u>	<u>A077438 001</u>	Jun 20, 2011
<u>AB</u>			<u>500MG</u>	<u>A077438 002</u>	Jun 20, 2011
<u>AB</u>			<u>750MG</u>	<u>A077438 003</u>	Jun 20, 2011
<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>		TORRENT PHARMS	<u>250MG</u>	<u>A090722 001</u>	Jun 20, 2011

PRESCRIPTION DRUG PRODUCT LIST

LEVOFLOXACIN

TABLET; ORAL

LEVOFLOXACIN

<u>AB</u>		<u>500MG</u>	<u>A090722</u>	<u>002</u>	Jun 20, 2011
<u>AB</u>		<u>750MG</u>	<u>A090722</u>	<u>003</u>	Jun 20, 2011
<u>AB</u>	ZYDUS PHARMS USA INC	<u>250MG</u>	<u>A077652</u>	<u>001</u>	Sep 07, 2012
<u>AB</u>		<u>500MG</u>	<u>A077652</u>	<u>002</u>	Sep 07, 2012
<u>AB</u>		<u>750MG</u>	<u>A077652</u>	<u>003</u>	Sep 07, 2012

LEVOLEUCOVORIN

POWDER; INTRAVENOUS

KHAPZORY

+	!	ACROTECH	175MG/VIAL	N211226	001	Oct 19, 2018
+	!		300MG/VIAL	N211226	002	Oct 19, 2018

LEVOLEUCOVORIN CALCIUM

POWDER; INTRAVENOUS

FUSILEV

<u>AP</u>	+	!	ACROTECH	<u>EQ 50MG BASE/VIAL</u>	<u>N020140</u>	<u>001</u>	Mar 07, 2008
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LEVOLEUCOVORIN CALCIUM

<u>AP</u>			ACTAVIS LLC	<u>EQ 50MG BASE/VIAL</u>	<u>A206516</u>	<u>001</u>	Feb 13, 2017
<u>AP</u>			AMNEAL	<u>EQ 50MG BASE/VIAL</u>	<u>A207547</u>	<u>001</u>	Feb 13, 2017
<u>AP</u>			MEITHEAL	<u>EQ 50MG BASE/VIAL</u>	<u>A211003</u>	<u>001</u>	Aug 22, 2019
<u>AP</u>			WEST-WARD PHARMS INT	<u>EQ 50MG BASE/VIAL</u>	<u>A206263</u>	<u>001</u>	Jun 16, 2016

SOLUTION; INTRAVENOUS

LEVOLEUCOVORIN CALCIUM

<u>AP</u>			AMNEAL	<u>EQ 175MG BASE/17.5ML (EQ 10MG BASE/ML)</u>	<u>A207548</u>	<u>001</u>	Sep 08, 2017
<u>AP</u>			GLAND PHARMA LTD	<u>EQ 175MG BASE/17.5ML (EQ 10MG BASE/ML)</u>	<u>A210892</u>	<u>001</u>	Sep 14, 2018
<u>AP</u>				<u>EQ 250MG BASE/25ML (EQ 10MG BASE/ML)</u>	<u>A210892</u>	<u>002</u>	Sep 14, 2018
<u>AP</u>			INGENUS PHARMS LLC	<u>EQ 175MG BASE/17.5ML (EQ 10MG BASE/ML)</u>	<u>A210623</u>	<u>001</u>	May 03, 2018
<u>AP</u>				<u>EQ 250MG BASE/25ML (EQ 10MG BASE/ML)</u>	<u>A210623</u>	<u>002</u>	May 03, 2018
<u>AP</u>			MEITHEAL	<u>EQ 175MG BASE/17.5ML (EQ 10MG BASE/ML)</u>	<u>A211002</u>	<u>001</u>	Aug 16, 2019
<u>AP</u>				<u>EQ 250MG BASE/25ML (EQ 10MG BASE/ML)</u>	<u>A211002</u>	<u>002</u>	Aug 16, 2019
<u>AP</u>			SANDOZ INC	<u>EQ 175MG BASE/17.5ML (EQ 10MG BASE/ML)</u>	<u>A203563</u>	<u>001</u>	Mar 09, 2015
<u>AP</u>			!	<u>EQ 250MG BASE/25ML (EQ 10MG BASE/ML)</u>	<u>A203563</u>	<u>002</u>	Mar 09, 2015

LEVOMILNACIPRAN HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

FETZIMA

+			ALLERGAN	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

LEVONORDEFIN; MEPIVACAINE HYDROCHLORIDE

INJECTABLE; INJECTION

SCANDONEST L

!			DEPROCO	0.05MG/ML; 2%	A088388	001	Oct 10, 1984
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LEVONORGESTREL

INTRAUTERINE DEVICE; INTRAUTERINE

KYLEENA

+	!		BAYER HLTHCARE	19.5MG	N208224	001	Sep 16, 2016
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LILETTA

			MEDICINES360	52MG	N206229	001	Feb 26, 2015
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MIRENA

+	!		BAYER HLTHCARE	52MG	N021225	001	Dec 06, 2000
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SKYLA

+	!		BAYER HLTHCARE	13.5MG	N203159	001	Jan 09, 2013
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TABLET; ORAL

LEVONORGESTREL

<u>AB</u>	!		L PERRIGO CO	<u>0.75MG</u>	<u>A090740</u>	<u>001</u>	Dec 30, 2010
<u>AB</u>			MYLAN LABS LTD	<u>0.75MG</u>	<u>A202740</u>	<u>001</u>	Sep 02, 2016

LEVORPHANOL TARTRATE

TABLET; ORAL

LEVORPHANOL TARTRATE

<u>AB</u>	!		SENTYNL THERAPS INC	<u>2MG</u>	<u>A074278</u>	<u>001</u>	Mar 31, 2000
<u>AB</u>			SPECGX LLC	<u>2MG</u>	<u>A212024</u>	<u>001</u>	Dec 13, 2019
<u>AB</u>			VIRTUS PHARMS	<u>2MG</u>	<u>A211484</u>	<u>001</u>	Dec 13, 2018
	!		SENTYNL THERAPS INC	3MG	A074278	003	Jun 18, 2018

PRESCRIPTION DRUG PRODUCT LIST

LEVOTHYROXINE SODIUM

CAPSULE; ORAL

TIROSINT

+	INSTITUT BIOCHIMIQUE	0.013MG	N021924 013	Aug 01, 2007
+		0.025MG	N021924 002	Oct 13, 2006
+		0.05MG	N021924 003	Oct 13, 2006
+		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
+		0.200MG	N021924 012	Apr 25, 2017

POWDER; INTRAVENOUS

LEVOTHYROXINE SODIUM

AP	+ !	FRESENIUS KABI USA	<u>100MCG/VIAL</u>	<u>N202231 001</u>	Jun 24, 2011
AP	+ !		<u>200MCG/VIAL</u>	<u>N202231 002</u>	Jun 24, 2011
AP	+ !		<u>500MCG/VIAL</u>	<u>N202231 003</u>	Jun 24, 2011
AP		MAIA PHARMS INC	<u>100MCG/VIAL</u>	<u>A208749 001</u>	Dec 21, 2018
AP			<u>200MCG/VIAL</u>	<u>A208749 002</u>	Dec 21, 2018
AP			<u>500MCG/VIAL</u>	<u>A208749 003</u>	Dec 21, 2018
AP		PIRAMAL CRITICAL	<u>100MCG/VIAL</u>	<u>A206163 001</u>	Jun 29, 2016
AP			<u>500MCG/VIAL</u>	<u>A206163 002</u>	Jun 29, 2016

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

SOLUTION; ORAL

TIROSINT-SOL

+	INSTITUT BIOCHIMIQUE	13MCG/ML	N206977 001	Dec 15, 2016
+		25MCG/ML	N206977 002	Dec 15, 2016
+		50MCG/ML	N206977 003	Dec 15, 2016
+		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

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
-->			--> <u>AB1, AB2, AB3</u>	<u>0.1MG</u>	N021342 005	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>0.112MG</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 ATLANTIS	--> <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

LEVOTHYROXINE SODIUM

-->	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>AB1,AB2,</u>	<u>0.112MG</u>	A076187 006	Jun 05, 2002

PRESCRIPTION DRUG PRODUCT LIST

LEVOTHYROXINE SODIUM **

**See current Annual Edition, 1.8 Description of Special Situations, Levothyroxine Sodium

TABLET;ORAL

LEVOTHYROXINE SODIUM

		<u>AB3,AB4</u>	<u>0.112MG</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>					

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</u>	<u>001</u>	May 31,	2002
<u>AB2</u>		<u>0.05MG</u>	<u>N021292</u>	<u>002</u>	May 31,	2002
<u>AB2</u>		<u>0.075MG</u>	<u>N021292</u>	<u>003</u>	May 31,	2002
<u>AB2</u>		<u>0.088MG</u>	<u>N021292</u>	<u>004</u>	May 31,	2002
<u>AB2</u>		<u>0.1MG</u>	<u>N021292</u>	<u>005</u>	May 31,	2002
<u>AB2</u>		<u>0.112MG</u>	<u>N021292</u>	<u>006</u>	May 31,	2002
<u>AB2</u>		<u>0.125MG</u>	<u>N021292</u>	<u>007</u>	May 31,	2002
<u>AB2</u>		<u>0.137MG</u>	<u>N021292</u>	<u>008</u>	May 31,	2002
<u>AB2</u>		<u>0.15MG</u>	<u>N021292</u>	<u>009</u>	May 31,	2002
<u>AB2</u>		<u>0.175MG</u>	<u>N021292</u>	<u>010</u>	May 31,	2002
<u>AB2</u>		<u>0.2MG</u>	<u>N021292</u>	<u>011</u>	May 31,	2002

THYRO-TABS

-->	+	ALVOGEN	-->	<u>AB2,AB4</u>	<u>0.025MG</u>	N021116	001	Oct 24,	2002
-->	+		-->	<u>AB2,AB4</u>	<u>0.05MG</u>	N021116	002	Oct 24,	2002
-->	+		-->	<u>AB2,AB4</u>	<u>0.075MG</u>	N021116	003	Oct 24,	2002
-->	+		-->	<u>AB2,AB4</u>	<u>0.088MG</u>	N021116	010	Oct 24,	2002
-->	+		-->	<u>AB2,AB4</u>	<u>0.1MG</u>	N021116	004	Oct 24,	2002
-->	+		-->	<u>AB2,AB4</u>	<u>0.112MG</u>	N021116	011	Oct 24,	2002
-->	+		-->	<u>AB2,AB4</u>	<u>0.125MG</u>	N021116	005	Oct 24,	2002
-->	+		-->	<u>AB2,AB4</u>	<u>0.137MG</u>	N021116	012	Dec 07,	2004
-->	+		-->	<u>AB2,AB4</u>	<u>0.15MG</u>	N021116	006	Oct 24,	2002
-->	+		-->	<u>AB2,AB4</u>	<u>0.175MG</u>	N021116	007	Oct 24,	2002
-->	+		-->	<u>AB2,AB4</u>	<u>0.2MG</u>	N021116	008	Oct 24,	2002
-->	+		-->	<u>AB2,AB4</u>	<u>0.3MG</u>	N021116	009	Oct 24,	2002

LIDOCAINE

OINTMENT;TOPICAL

LIDOCAINE

<u>AT</u>	ACP NIMBLE	<u>5%</u>	<u>A211019</u>	<u>001</u>	Dec 12,	2018
<u>AT</u>	ALEOR	<u>5%</u>	<u>A211469</u>	<u>001</u>	Nov 23,	2018
<u>AT</u>	DERMACEUTICALS					
<u>AT</u>	ALKEM LABS LTD	<u>5%</u>	<u>A207810</u>	<u>001</u>	Mar 10,	2017
<u>AT</u>	AMNEAL PHARMS	<u>5%</u>	<u>A206297</u>	<u>001</u>	Aug 07,	2015
<u>AT</u>	!	FOUGERA PHARMS INC	<u>5%</u>	<u>A080198</u>	<u>001</u>	
<u>AT</u>	GLENMARK PHARMS LTD	<u>5%</u>	<u>A206498</u>	<u>001</u>	Sep 09,	2016
<u>AT</u>	MACLEODS PHARMS LTD	<u>5%</u>	<u>A211697</u>	<u>001</u>	Mar 16,	2020
<u>AT</u>	NOVAST LABS	<u>5%</u>	<u>A208604</u>	<u>001</u>	Sep 20,	2017
<u>AT</u>	SEPTODONT INC	<u>5%</u>	<u>A040911</u>	<u>001</u>	May 23,	2011
<u>AT</u>	STRIDES PHARMA	<u>5%</u>	<u>A210958</u>	<u>001</u>	Dec 11,	2018
<u>AT</u>	SUNGEN PHARMA	<u>5%</u>	<u>A212486</u>	<u>001</u>	Oct 17,	2019
<u>AT</u>	TARO	<u>5%</u>	<u>A086724</u>	<u>001</u>		
<u>AT</u>	TELIGENT PHARMA INC	<u>5%</u>	<u>A205318</u>	<u>001</u>	Feb 01,	2016
<u>AT</u>	TEVA PHARMS USA	<u>5%</u>	<u>A210256</u>	<u>001</u>	Jan 16,	2018
<u>AT</u>	VITRUVIAS THERAP	<u>5%</u>	<u>A208822</u>	<u>001</u>	Sep 25,	2017

PRESCRIPTION DRUG PRODUCT LIST

LIDOCAINE

PATCH; TOPICAL

LIDOCAINE

AB	ACTAVIS LABS UT INC	5%	A200675	001	Aug 23, 2012
AB	MYLAN TECHNOLOGIES	5%	A202346	001	Aug 07, 2015

LIDODERM

AB	+ !	TEIKOKU PHARMA USA	5%	N020612	001	Mar 19, 1999
		ZTLIDO				
	+ !	SCILEX PHARMS INC	1.8%	N207962	001	Feb 28, 2018

LIDOCAINE HYDROCHLORIDE

GEL; OPHTHALMIC

AKTEN

+ !	AKORN	3.5%	N022221	001	Oct 07, 2008
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INJECTABLE; INJECTION

LIDOCAINE HYDROCHLORIDE

AP	AUROBINDO PHARMA LTD	1%	A207182	001	Oct 30, 2017
AP		2%	A207182	002	Oct 30, 2017
AP	B BRAUN MEDICAL INC	1%	A208474	001	Aug 03, 2018
AP	FRESENIUS KABI USA	1%	A080404	002	
AP		2%	A080404	003	
AP	HOSPIRA	0.5%	A088328	001	May 17, 1984
AP	!	1%	A083158	001	
AP		1%	A088329	001	May 17, 1984
AP		2%	A040078	001	Jun 23, 1995
AP	!	2%	A083158	002	
AP		2%	A088294	001	May 17, 1984
AP	INTL MEDICATION	1%	A083173	001	
AP		2%	A083173	002	
AP	SPECTRA MDCL DEVICES	1%	A208017	001	Apr 18, 2018

LIDOCAINE HYDROCHLORIDE 0.2% AND DEXTROSE 5% IN PLASTIC CONTAINER

AP	B BRAUN	200MG/100ML	N019830	002	Apr 08, 1992
AP	BAXTER HLTHCARE	200MG/100ML	N018461	002	

LIDOCAINE HYDROCHLORIDE 0.4% AND DEXTROSE 5% IN PLASTIC CONTAINER

AP	B BRAUN	400MG/100ML	N019830	003	Apr 08, 1992
AP	BAXTER HLTHCARE	400MG/100ML	N018461	003	

LIDOCAINE HYDROCHLORIDE 0.8% AND DEXTROSE 5% IN PLASTIC CONTAINER

AP	B BRAUN	800MG/100ML	N019830	004	Apr 08, 1992
AP	BAXTER HLTHCARE	800MG/100ML	N018461	004	Feb 22, 1982

LIDOCAINE HYDROCHLORIDE IN PLASTIC CONTAINER

AP	FRESENIUS KABI USA	1%	A088586	001	Jul 24, 1985
AP	HOSPIRA	0.5%	A088325	001	Jul 31, 1984
AP		1%	A088299	001	Jul 31, 1984
AP		2%	A088327	001	Jul 31, 1984

LIDOCAINE HYDROCHLORIDE PRESERVATIVE FREE

AP	AUROBINDO PHARMA LTD	1%	A203040	001	Mar 14, 2013
AP		1%	A203082	001	Mar 14, 2013
AP		2%	A203040	002	Mar 14, 2013
AP		2%	A203082	002	Mar 14, 2013
AP	FRESENIUS KABI USA	2%	N017584	001	
AP		4%	N017584	002	
AP	HOSPIRA	1%	A080408	001	
AP		1.5%	A080408	002	

LIDOCAINE HYDROCHLORIDE PRESERVATIVE FREE IN PLASTIC CONTAINER

AP	HOSPIRA	1%	A040302	001	Sep 28, 1998
AP		2%	A040302	002	Sep 28, 1998

XYLOCAINE

AP	+ !	FRESENIUS KABI USA	0.5%	N006488	008
AP	+ !		1%	N006488	007
AP	+ !		1.5%	N006488	010
AP	+ !		2%	N006488	002

LIDOCAINE HYDROCHLORIDE PRESERVATIVE FREE

!	HOSPIRA	4%	A088295	001	May 17, 1984
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INJECTABLE; SPINAL

LIDOCAINE HYDROCHLORIDE 5% AND DEXTROSE 7.5%

!	HOSPIRA	5%	A083914	001	
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JELLY; TOPICAL

GLYDO

AT	SAGENT PHARMS INC	2%	A201094	001	Apr 28, 2014
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PRESCRIPTION DRUG PRODUCT LIST

LIDOCAINE HYDROCHLORIDE

JELLY; TOPICAL

LIDOCAINE HYDROCHLORIDE

AT	AKORN	2%	A040433 001	Feb 12, 2003
AT	INTL MEDICATION	2%	A086283 001	

XYLOCAINE

AT	+! OAK PHARMS	2%	N008816 001	
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SOLUTION; ORAL

LIDOCAINE HYDROCHLORIDE

AT	HI TECH PHARMA	2%	A040014 001	Jul 10, 1995
AT	WOCKHARDT BIO AG	2%	A087872 001	Nov 18, 1982

LIDOCAINE HYDROCHLORIDE VISCOUS

AT	! LANNETT CO INC	2%	A040708 001	Feb 27, 2007
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LIDOCAINE VISCOUS

AT	HIKMA	2%	A088802 001	Apr 26, 1985
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SOLUTION; TOPICAL

LARYNG-O-JET KIT

AT	INTL MEDICATION	4%	A086364 001	
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LIDOCAINE HYDROCHLORIDE

AT	! HIKMA	4%	A088803 001	Apr 03, 1985
AT	LANNETT CO INC	4%	A040710 001	Feb 27, 2007
AT	TELOGENT PHARMA INC	4%	A204494 001	Mar 12, 2014

SYSTEM; INTRADERMAL

ZINGO

POWDER PHARMS	0.5MG		N022114 001	Aug 16, 2007
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LIDOCAINE; PRILUCAINE

CREAM; TOPICAL

EMLA

AB	+! TEVA BRANDED PHARM	2.5%;2.5%	N019941 001	Dec 30, 1992
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LIDOCAINE AND PRILUCAINE

AB	FOUGERA PHARMS	2.5%;2.5%	A076453 001	Aug 18, 2003
AB	HI TECH PHARMA	2.5%;2.5%	A076290 001	Sep 25, 2003
AB	TELOGENT PHARMA INC	2.5%;2.5%	A205887 001	Jun 29, 2018
AB	TOLMAR	2.5%;2.5%	A076320 001	Aug 27, 2003

GEL; PERIODONTAL

ORAQIX

+! DENTSPLY PHARM	2.5%;2.5%		N021451 001	Dec 19, 2003
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LIDOCAINE; TETRACAINE

CREAM; TOPICAL

PLIAGLIS

+! TARO PHARMS	7%;7%		N021717 001	Jun 29, 2006
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PATCH; TOPICAL

SYNERA

+! GALEN SPECIALTY	70MG;70MG		N021623 001	Jun 23, 2005
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LIFITEGRAST

SOLUTION/DROPS; OPHTHALMIC

XIIDRA

+! NOVARTIS	5%		N208073 001	Jul 11, 2016
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LINACLOTIDE

CAPSULE; ORAL

LINZESS

+! ALLERGAN	72MCG		N202811 003	Jan 25, 2017
+!	145MCG		N202811 001	Aug 30, 2012
+	290MCG		N202811 002	Aug 30, 2012

LINAGLIPTIN

TABLET; ORAL

TRADJENTA

+! BOEHRINGER INGELHEIM	5MG		N201280 001	May 02, 2011
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LINAGLIPTIN; METFORMIN HYDROCHLORIDE

TABLET; ORAL

JENTADUETO

+ BOEHRINGER INGELHEIM	2.5MG;500MG		N201281 001	Jan 30, 2012
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+	2.5MG;850MG		N201281 002	Jan 30, 2012
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+!	2.5MG;1GM		N201281 003	Jan 30, 2012
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TABLET, EXTENDED RELEASE; ORAL

JENTADUETO XR

+ BOEHRINGER INGELHEIM	2.5MG;1GM		N208026 001	May 27, 2016
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PRESCRIPTION DRUG PRODUCT LIST

LINAGLIPTIN; METFORMIN HYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL

JENTADUETO XR

+!

5MG;1GM

N208026 002 May 27, 2016

LINCOMYCIN HYDROCHLORIDE

INJECTABLE; INJECTION

LINCOCINAP +! PHARMACIA AND
UPJOHNEQ 300MG BASE/MLN050317 001LINCOMYCINAP XGEN PHARMSEQ 300MG BASE/MLA201746 001 Jun 04, 2015LINDANE

LOTION; TOPICAL

LINDANE

! OLTA PHARMS

1%

A087313 001

SHAMPOO; TOPICAL

LINDANEAT OLTA PHARMS1%A087266 001AT ! WOCKHARDT BIO AG1%A088191 001 Sep 18, 1984LINEZOLID

FOR SUSPENSION; ORAL

LINEZOLIDAB HIKMA100MG/5MLA200068 001 Jun 03, 2015ZYVOXAB +! PHARMACIA AND
UPJOHN100MG/5MLN021132 001 Apr 18, 2000

SOLUTION; INTRAVENOUS

LINEZOLIDAP AUROBINDO PHARMA
LTD600MG/300ML (2MG/ML)A206917 001 Aug 04, 2016AP FRESENIUS KABI USA600MG/300ML (2MG/ML)A204764 001 Mar 15, 2016AP HOSPIRA INC600MG/300ML (2MG/ML)A205442 001 Jul 07, 2015AP HQ SPCLT PHARMA200MG/100ML (2MG/ML)A207001 001 Jul 07, 2017AP600MG/300ML (2MG/ML)A207001 002 Jul 07, 2017AP MYLAN LABS LTD200MG/100ML (2MG/ML)A205154 001 Dec 06, 2017AP600MG/300ML (2MG/ML)A205154 002 Dec 06, 2017AP NANG KUANG PHARM CO200MG/100ML (2MG/ML)A207354 001 Dec 20, 2016AP600MG/300ML (2MG/ML)A207354 002 Dec 20, 2016AP SAGENT PHARMS INC200MG/100ML (2MG/ML)A204696 001 Mar 02, 2017AP600MG/300ML (2MG/ML)A204696 002 Mar 02, 2017AP SANDOZ INC200MG/100ML (2MG/ML)A200904 001 Jul 16, 2015AP600MG/300ML (2MG/ML)A200904 002 Jul 16, 2015AP TEVA PHARMS600MG/300ML (2MG/ML)A200222 001 Jun 27, 2012ZYVOXAP + PHARMACIA AND
UPJOHN200MG/100ML (2MG/ML)N021131 001 Apr 18, 2000AP +!600MG/300ML (2MG/ML)N021131 003 Apr 18, 2000

LINEZOLID IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

+! HOSPIRA INC

600MG/300ML (2MG/ML)

N206473 001 Jun 18, 2015

TABLET; ORAL

LINEZOLIDAB ALEMBIC PHARMS LTD600MGA205233 001 Dec 21, 2015AB ALKEM LABS LTD600MGA205517 001 Dec 21, 2015AB AMNEAL PHARMS600MGA204536 001 Dec 21, 2015AB CELLTRION600MGA210702 001 Apr 25, 2019AB GATE PHARMS600MGA091210 001 Feb 05, 2016AB GLENMARK PHARMS600MGA078987 001 Dec 21, 2015AB HETERO LABS LTD V600MGA204239 001 Dec 21, 2015AB NOVEL LABS INC600MGA207526 001 Aug 22, 2016AB TEVA PHARMS USA600MGA078061 001 May 18, 2015AB ZYDUS PHARMS600MGA206097 001 Feb 22, 2017ZYVOXAB +! PHARMACIA AND
UPJOHN600MGN021130 002 Apr 18, 2000LIOTHYRONINE SODIUM

INJECTABLE; INJECTION

LIOTHYRONINE SODIUMAP XGEN PHARMSEQ 0.01MG BASE/MLA076923 001 Aug 17, 2005TRIOSTATAP +! PAR STERILE
PRODUCTSEQ 0.01MG BASE/MLN020105 001 Dec 31, 1991

PRESCRIPTION DRUG PRODUCT LIST

3-269 (of 453)

LIOTHYRONINE SODIUM

TABLET; ORAL

CYTOMEL

<u>AB</u>	+	KING PHARMS	<u>EQ 0.005MG BASE</u>	<u>N010379</u>	<u>001</u>	
<u>AB</u>	+		<u>EQ 0.025MG BASE</u>	<u>N010379</u>	<u>002</u>	
<u>AB</u>	+		<u>EQ 0.05MG BASE</u>	<u>N010379</u>	<u>003</u>	

LIOTHYRONINE SODIUM

<u>AB</u>		MAYNE PHARMA INC	<u>EQ 0.005MG BASE</u>	<u>A090097</u>	<u>001</u>	Mar 20, 2009
<u>AB</u>			<u>EQ 0.025MG BASE</u>	<u>A090097</u>	<u>002</u>	Mar 20, 2009
<u>AB</u>			<u>EQ 0.05MG BASE</u>	<u>A090097</u>	<u>003</u>	Mar 20, 2009
<u>AB</u>		SIGMAPHARM LABS LLC	<u>EQ 0.005MG BASE</u>	<u>A200295</u>	<u>001</u>	Nov 29, 2012
<u>AB</u>			<u>EQ 0.025MG BASE</u>	<u>A200295</u>	<u>002</u>	Nov 29, 2012
<u>AB</u>	!		<u>EQ 0.05MG BASE</u>	<u>A200295</u>	<u>003</u>	Nov 29, 2012
<u>AB</u>		SUN PHARM	<u>EQ 0.005MG BASE</u>	<u>A091382</u>	<u>001</u>	Apr 20, 2016
<u>AB</u>			<u>EQ 0.025MG BASE</u>	<u>A091382</u>	<u>002</u>	Apr 20, 2016
<u>AB</u>			<u>EQ 0.05MG BASE</u>	<u>A091382</u>	<u>003</u>	Apr 20, 2016

LIRAGLUTIDE RECOMBINANT

SOLUTION; SUBCUTANEOUS

SAXENDA

+	!	NOVO	18MG/3ML (6MG/ML)	N206321	001	Dec 23, 2014
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VICTOZA

+	!	NOVO NORDISK INC	18MG/3ML (6MG/ML)	N022341	001	Jan 25, 2010
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LISDEXAMFETAMINE DIMESYLATE

CAPSULE; ORAL

VYVANSE

+		SHIRE DEVELOPMENT	10MG	N021977	007	Oct 30, 2014
+			20MG	N021977	004	Dec 10, 2007
+			30MG	N021977	001	Feb 23, 2007
+			40MG	N021977	005	Dec 10, 2007
+			50MG	N021977	002	Feb 23, 2007
+			60MG	N021977	006	Dec 10, 2007
+	!		70MG	N021977	003	Feb 23, 2007

TABLET, CHEWABLE; ORAL

VYVANSE

+		SHIRE DEV LLC	10MG	N208510	001	Jan 28, 2017
+			20MG	N208510	002	Jan 28, 2017
+			30MG	N208510	003	Jan 28, 2017
+			40MG	N208510	004	Jan 28, 2017
+			50MG	N208510	005	Jan 28, 2017
+	!		60MG	N208510	006	Jan 28, 2017

LISINAPRIL

SOLUTION; ORAL

QBRELIS

+	!	SILVERGATE PHARMS	1MG/ML	N208401	001	Jul 29, 2016
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TABLET; ORAL

LISINAPRIL

<u>AB</u>		ACCORD HLTHCARE	<u>2.5MG</u>	<u>A202554</u>	<u>001</u>	Jul 30, 2013
<u>AB</u>			<u>5MG</u>	<u>A202554</u>	<u>002</u>	Jul 30, 2013
<u>AB</u>			<u>10MG</u>	<u>A202554</u>	<u>003</u>	Jul 30, 2013
<u>AB</u>			<u>20MG</u>	<u>A202554</u>	<u>004</u>	Jul 30, 2013
<u>AB</u>			<u>30MG</u>	<u>A202554</u>	<u>005</u>	Jul 30, 2013
<u>AB</u>			<u>40MG</u>	<u>A202554</u>	<u>006</u>	Jul 30, 2013
<u>AB</u>		AUROBINDO	<u>2.5MG</u>	<u>A077622</u>	<u>001</u>	Feb 22, 2006
<u>AB</u>			<u>5MG</u>	<u>A077622</u>	<u>002</u>	Feb 22, 2006
<u>AB</u>			<u>10MG</u>	<u>A077622</u>	<u>003</u>	Feb 22, 2006
<u>AB</u>			<u>20MG</u>	<u>A077622</u>	<u>004</u>	Feb 22, 2006
<u>AB</u>			<u>30MG</u>	<u>A077622</u>	<u>005</u>	Feb 22, 2006
<u>AB</u>			<u>40MG</u>	<u>A077622</u>	<u>006</u>	Feb 22, 2006
<u>AB</u>		CASI PHARMS INC	<u>2.5MG</u>	<u>A075994</u>	<u>001</u>	Jul 01, 2002
<u>AB</u>			<u>5MG</u>	<u>A075994</u>	<u>002</u>	Jul 01, 2002
<u>AB</u>			<u>10MG</u>	<u>A075994</u>	<u>003</u>	Jul 01, 2002
<u>AB</u>			<u>20MG</u>	<u>A075994</u>	<u>004</u>	Jul 01, 2002
<u>AB</u>			<u>30MG</u>	<u>A075994</u>	<u>005</u>	Jul 01, 2002
<u>AB</u>			<u>40MG</u>	<u>A075994</u>	<u>006</u>	Jul 01, 2002
<u>AB</u>		COREPHARMA	<u>2.5MG</u>	<u>A076102</u>	<u>001</u>	Sep 30, 2002
<u>AB</u>			<u>5MG</u>	<u>A076102</u>	<u>002</u>	Sep 30, 2002
<u>AB</u>			<u>10MG</u>	<u>A076102</u>	<u>003</u>	Sep 30, 2002
<u>AB</u>			<u>20MG</u>	<u>A076102</u>	<u>004</u>	Sep 30, 2002
<u>AB</u>			<u>30MG</u>	<u>A076102</u>	<u>005</u>	Sep 30, 2002
<u>AB</u>			<u>40MG</u>	<u>A076102</u>	<u>006</u>	Sep 30, 2002
<u>AB</u>		INVAGEN PHARMS	<u>2.5MG</u>	<u>A203508</u>	<u>001</u>	Oct 29, 2013

PRESCRIPTION DRUG PRODUCT LIST

LISINAPRIL

TABLET; ORAL

LISINAPRIL

<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>A076180 001</u>	Jul 01, 2002
<u>AB</u>		<u>5MG</u>	<u>A075743 002</u>	Jul 01, 2002
<u>AB</u>		<u>5MG</u>	<u>A076180 002</u>	Jul 01, 2002
<u>AB</u>		<u>10MG</u>	<u>A075743 003</u>	Jul 01, 2002
<u>AB</u>		<u>10MG</u>	<u>A076180 003</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>	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	<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

PRINIVIL

<u>AB</u>	MERCK	<u>5MG</u>	<u>N019558 001</u>	Dec 29, 1987
<u>AB</u>		<u>40MG</u>	<u>N019558 004</u>	Oct 25, 1988

ZESTRIL

<u>AB</u>	+ ALVOGEN	<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 GENERICS	<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

PRESCRIPTION DRUG PRODUCT LISTLITHIUM CARBONATE

TABLET; ORAL

LITHIUM CARBONATE

AB	+ !	HIKMA	300MG	N018558 001	Jan 29, 1982
AB		SUN PHARM INDS INC	300MG	A091027 001	Jun 24, 2010

TABLET, EXTENDED RELEASE; ORAL

LITHIUM CARBONATE

AB		ALEMBIC PHARMS LTD	300MG	A204445 001	Jun 10, 2015
AB		GLENMARK GENERICS	450MG	A091616 001	Feb 14, 2011
AB		GLENMARK PHARMS INC	300MG	A091544 001	Dec 27, 2010
AB		HERITAGE PHARMA	300MG	A205532 001	Sep 29, 2016
AB		HIKMA	300MG	A076832 001	Oct 28, 2004
AB	!		450MG	A076691 001	Jan 05, 2004
AB		MYLAN PHARMS INC	300MG	A202288 001	Jun 29, 2012
AB			450MG	A202219 001	Aug 08, 2012
AB		UNIQUE PHARM LABS	300MG	A204779 001	Jul 27, 2015
AB			450MG	A205663 001	Jun 05, 2017

LITHIOBID

AB	+ !	ANI PHARMS INC	300MG	N018027 001	
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LITHIUM CITRATE

SYRUP; ORAL

LITHIUM CITRATE

AA	+ !	HIKMA	EQ 300MG CARBONATE/5ML	N018421 001	
AA		WOCKHARDT BIO AG	EQ 300MG CARBONATE/5ML	A070755 001	May 21, 1986

LIXISENATIDE

SOLUTION; SUBCUTANEOUS

ADLYXIN

	+ !	SANOFI-AVENTIS US	0.15MG/3ML (0.05MG/ML)	N208471 001	Jul 27, 2016
	+ !		0.3MG/3ML (0.1MG/ML)	N208471 002	Jul 27, 2016

LODOXAMIDE TROMETHAMINE

SOLUTION/DROPS; OPHTHALMIC

ALOMIDE

	+ !	NOVARTIS	EQ 0.1% BASE	N020191 001	Sep 23, 1993
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LOFEXIDINE HYDROCHLORIDE

TABLET; ORAL

LUCEMYRA

	+ !	US WORLDMEDS LLC	EQ 0.18MG BASE	N209229 001	May 16, 2018
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LOMITAPIDE MESYLATE

CAPSULE; ORAL

JUXTAPID

	+	AEGERION	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
	+		EQ 40MG BASE	N203858 005	Apr 23, 2015
	+ !		EQ 60MG BASE	N203858 006	Apr 23, 2015

LOMUSTINE

CAPSULE; ORAL

GLEOSTINE

	+	CORDEN PHARMA	10MG	N017588 001	
	+		40MG	N017588 002	
	+ !		100MG	N017588 003	

LOPERAMIDE HYDROCHLORIDE

CAPSULE; ORAL

LOPERAMIDE HYDROCHLORIDE

AB	!	MYLAN	2MG	A072741 001	Sep 18, 1991
AB		TEVA	2MG	A073192 001	Apr 30, 1992

LOPINAVIR; RITONAVIR

SOLUTION; ORAL

KALETRA

AA	+ !	ABBVIE	80MG/ML; 20MG/ML	N021251 001	Sep 15, 2000
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LOPINAVIR AND RITONAVIR

AA		LANNETT CO INC	80MG/ML; 20MG/ML	A207407 001	Dec 27, 2016
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TABLET; ORAL

KALETRA

	+	ABBVIE	100MG; 25MG	N021906 002	Nov 09, 2007
	+ !		200MG; 50MG	N021906 001	Oct 28, 2005

PRESCRIPTION DRUG PRODUCT LIST

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LORAZEPAM

CONCENTRATE; ORAL

LORAZEPAM

<u>AA</u>	AMNEAL PHARMS	<u>2MG/ML</u>	<u>A091383</u>	<u>001</u>	Dec 23, 2009
<u>AA</u>	HL-TECH PHARMA CO	<u>2MG/ML</u>	<u>A200169</u>	<u>001</u>	Jan 30, 2012
<u>AA</u>	LANNETT CO INC	<u>2MG/ML</u>	<u>A079244</u>	<u>001</u>	Apr 28, 2009
<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
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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>	AKORN	<u>2MG/ML</u>	<u>A075025</u>	<u>001</u>	Jul 23, 1998
<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

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>	AMNEAL PHARMS	<u>0.5MG</u>	<u>A078826</u>	<u>001</u>	Jun 23, 2010
<u>AB</u>		<u>1MG</u>	<u>A078826</u>	<u>002</u>	Jun 23, 2010
<u>AB</u>		<u>2MG</u>	<u>A078826</u>	<u>003</u>	Jun 23, 2010
<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 PHARMA LLC	<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>	MYLAN	<u>0.5MG</u>	<u>A077657</u>	<u>001</u>	Mar 16, 2006
<u>AB</u>		<u>1MG</u>	<u>A077657</u>	<u>002</u>	Mar 16, 2006
<u>AB</u>		<u>2MG</u>	<u>A077657</u>	<u>003</u>	Mar 16, 2006
<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>	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 INC	25MG	N210868	001	Nov 02, 2018
	+!	100MG	N210868	002	Nov 02, 2018

LOSARTAN POTASSIUM

TABLET; ORAL

COZAAR

<u>AB</u>	+! MERCK SHARP DOHME	<u>100MG</u>	<u>N020386</u>	<u>003</u>	Oct 13, 1998
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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>	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>	CADISTA PHARMS	<u>25MG</u>	<u>A201170</u>	<u>001</u>	Sep 18, 2012
<u>AB</u>		<u>50MG</u>	<u>A201170</u>	<u>002</u>	Sep 18, 2012
<u>AB</u>		<u>100MG</u>	<u>A201170</u>	<u>003</u>	Sep 18, 2012
<u>AB</u>	HETERO LABS LTD V	<u>25MG</u>	<u>A203835</u>	<u>001</u>	Aug 12, 2015
<u>AB</u>		<u>50MG</u>	<u>A203835</u>	<u>002</u>	Aug 12, 2015
<u>AB</u>		<u>100MG</u>	<u>A203835</u>	<u>003</u>	Aug 12, 2015
<u>AB</u>	IPCA LABS LTD	<u>25MG</u>	<u>A200290</u>	<u>001</u>	Aug 30, 2013

PRESCRIPTION DRUG PRODUCT LIST

3-273 (of 453)

LOSARTAN POTASSIUM

TABLET; ORAL

LOSARTAN POTASSIUM

<u>AB</u>		<u>50MG</u>	<u>A200290</u>	<u>002</u>	Aug 30, 2013
<u>AB</u>		<u>100MG</u>	<u>A200290</u>	<u>003</u>	Aug 30, 2013
<u>AB</u>	LUPIN LTD	<u>25MG</u>	<u>A078232</u>	<u>001</u>	Oct 06, 2010
<u>AB</u>		<u>50MG</u>	<u>A078232</u>	<u>002</u>	Oct 06, 2010
<u>AB</u>		<u>100MG</u>	<u>A078232</u>	<u>003</u>	Oct 06, 2010
<u>AB</u>	MACLEODS PHARMS LTD	<u>25MG</u>	<u>A202230</u>	<u>001</u>	May 30, 2012
<u>AB</u>		<u>50MG</u>	<u>A202230</u>	<u>002</u>	May 30, 2012
<u>AB</u>		<u>100MG</u>	<u>A202230</u>	<u>003</u>	May 30, 2012
<u>AB</u>	MICRO LABS	<u>25MG</u>	<u>A091541</u>	<u>001</u>	Sep 24, 2012
<u>AB</u>		<u>50MG</u>	<u>A091541</u>	<u>002</u>	Sep 24, 2012
<u>AB</u>		<u>100MG</u>	<u>A091541</u>	<u>003</u>	Sep 24, 2012
<u>AB</u>	MYLAN	<u>25MG</u>	<u>A091590</u>	<u>001</u>	Oct 06, 2010
<u>AB</u>		<u>50MG</u>	<u>A091590</u>	<u>002</u>	Oct 06, 2010
<u>AB</u>		<u>100MG</u>	<u>A091590</u>	<u>003</u>	Oct 06, 2010
<u>AB</u>	PRINSTON INC	<u>25MG</u>	<u>A091497</u>	<u>001</u>	Jun 06, 2011
<u>AB</u>		<u>50MG</u>	<u>A091497</u>	<u>002</u>	Jun 06, 2011
<u>AB</u>		<u>100MG</u>	<u>A091497</u>	<u>003</u>	Jun 06, 2011
<u>AB</u>	SANDOZ	<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>	STRIDES PHARMA	<u>25MG</u>	<u>A090382</u>	<u>001</u>	Oct 06, 2010
<u>AB</u>		<u>50MG</u>	<u>A090382</u>	<u>002</u>	Oct 06, 2010
<u>AB</u>		<u>100MG</u>	<u>A090382</u>	<u>003</u>	Oct 06, 2010
<u>AB</u>	TORRENT PHARMS	<u>25MG</u>	<u>A090467</u>	<u>001</u>	Oct 06, 2010
<u>AB</u>		<u>50MG</u>	<u>A090467</u>	<u>002</u>	Oct 06, 2010
<u>AB</u>		<u>100MG</u>	<u>A090467</u>	<u>003</u>	Oct 06, 2010
<u>AB</u>	UNICHEM LABS LTD	<u>25MG</u>	<u>A203030</u>	<u>001</u>	Oct 14, 2015
<u>AB</u>		<u>50MG</u>	<u>A203030</u>	<u>002</u>	Oct 14, 2015
<u>AB</u>		<u>100MG</u>	<u>A203030</u>	<u>003</u>	Oct 14, 2015
<u>AB</u>	UPSHER SMITH LABS	<u>25MG</u>	<u>A090544</u>	<u>001</u>	Oct 06, 2010
<u>AB</u>		<u>50MG</u>	<u>A090544</u>	<u>002</u>	Oct 06, 2010
<u>AB</u>		<u>100MG</u>	<u>A090544</u>	<u>003</u>	Oct 06, 2010
<u>AB</u>	WATSON LABS	<u>25MG</u>	<u>A091129</u>	<u>001</u>	Oct 06, 2010
<u>AB</u>		<u>50MG</u>	<u>A091129</u>	<u>002</u>	Oct 06, 2010
<u>AB</u>		<u>100MG</u>	<u>A091129</u>	<u>003</u>	Oct 06, 2010
<u>AB</u>	ZYDUS PHARMS USA INC	<u>25MG</u>	<u>A078243</u>	<u>001</u>	Oct 06, 2010
<u>AB</u>		<u>50MG</u>	<u>A078243</u>	<u>002</u>	Oct 06, 2010
<u>AB</u>		<u>100MG</u>	<u>A078243</u>	<u>003</u>	Oct 06, 2010

LOTEPREDNOL ETABONATE

GEL; OPTHALMIC

LOTEMAX

+! BAUSCH AND LOMB INC 0.5% N202872 001 Sep 28, 2012

LOTEMAX SM

+! BAUSCH AND LOMB INC 0.38% N208219 001 Feb 22, 2019

OINTMENT; OPTHALMIC

LOTEMAX

+! BAUSCH AND LOMB 0.5% N200738 001 Apr 15, 2011

SUSPENSION/DROPS; OPTHALMIC

LOTEMAXAB +! BAUSCH AND LOMB 0.5% N020583 001 Mar 09, 1998LOTEPREDNOL ETABONATEAB HI TECH 0.5% A207609 001 Apr 17, 2019

ALREX

+! BAUSCH AND LOMB 0.2% N020803 001 Mar 09, 1998

INVELTYS

+! KALA PHARMS INC 1% N210565 001 Aug 22, 2018

LOTEPREDNOL ETABONATE; TOBRAMYCIN

SUSPENSION/DROPS; OPTHALMIC

ZYLET

+! BAUSCH AND LOMB 0.5%;0.3% N050804 001 Dec 14, 2004

LOVASTATIN

TABLET; ORAL

LOVASTATIN

<u>AB</u>	ACTAVIS ELIZABETH	<u>10MG</u>	<u>A075828</u>	<u>001</u>	Dec 17, 2001
<u>AB</u>		<u>20MG</u>	<u>A075828</u>	<u>002</u>	Dec 17, 2001
<u>AB</u>		<u>40MG</u>	<u>A075828</u>	<u>003</u>	Dec 17, 2001
<u>AB</u>	BEXIMCO PHARMS USA	<u>10MG</u>	<u>A075300</u>	<u>001</u>	Dec 17, 2001

PRESCRIPTION DRUG PRODUCT LISTLOVASTATIN

TABLET; ORAL

LOVASTATIN

AB		20MG	A075300 002	Dec 17, 2001
AB		40MG	A075300 003	Dec 17, 2001
AB	CARLSBAD	10MG	A075991 001	Jun 05, 2002
AB		20MG	A075991 002	Jun 05, 2002
AB	!	40MG	A075991 003	Jun 05, 2002
AB	COREPHARMA	10MG	A077748 001	Feb 28, 2007
AB		20MG	A077748 002	Feb 28, 2007
AB		40MG	A077748 003	Feb 28, 2007
AB	LUPIN	10MG	A078296 001	Mar 14, 2008
AB		20MG	A078296 002	Nov 01, 2007
AB		40MG	A078296 003	Nov 01, 2007
AB	SANDOZ	10MG	A075636 001	Dec 17, 2001
AB		20MG	A075636 002	Dec 17, 2001
AB		40MG	A075636 003	Dec 17, 2001
AB	TEVA	10MG	A075551 003	Dec 17, 2001
AB		20MG	A075551 002	Dec 17, 2001
AB		40MG	A075551 001	Dec 17, 2001

TABLET, EXTENDED RELEASE; ORAL

ALTOPREV

+	COVIS PHARMA BV	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
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LOXAPINE SUCCINATE

CAPSULE; ORAL

LOXAPINE SUCCINATE

AB	ELITE LABS INC	EQ 5MG BASE	A076868 001	Aug 04, 2005
AB		EQ 10MG BASE	A076868 002	Aug 04, 2005
AB		EQ 25MG BASE	A076868 003	Aug 04, 2005
AB		EQ 50MG BASE	A076868 004	Aug 04, 2005
AB	LANNETT CO INC	EQ 5MG BASE	A090695 001	Sep 26, 2011
AB		EQ 10MG BASE	A090695 002	Sep 26, 2011
AB		EQ 25MG BASE	A090695 003	Sep 26, 2011
AB		EQ 50MG BASE	A090695 004	Sep 26, 2011
AB	MYLAN	EQ 5MG BASE	A076762 001	Nov 01, 2004
AB		EQ 10MG BASE	A076762 002	Nov 01, 2004
AB		EQ 25MG BASE	A076762 003	Nov 01, 2004
AB		EQ 50MG BASE	A076762 004	Nov 01, 2004
AB	WATSON LABS	EQ 5MG BASE	A072204 001	Jun 15, 1988
AB		EQ 10MG BASE	A072205 001	Jun 15, 1988
AB	!	EQ 25MG BASE	A072206 001	Jun 15, 1988
AB		EQ 50MG BASE	A072062 001	Jun 15, 1988

LUBIPROSTONE

CAPSULE; ORAL

AMITIZA

+	SUCAMPO PHARMA LLC	8MCG	N021908 002	Apr 29, 2008
+	!	24MCG	N021908 001	Jan 31, 2006

LULICONAZOLE

CREAM; TOPICAL

LUZU

+	MEDICIS	1%	N204153 001	Nov 14, 2013
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LUMATEPERONE TOSYLATE

CAPSULE; ORAL

CAPLYTA

+	INTRA-CELLULAR	EQ 42MG BASE	N209500 001	Dec 20, 2019
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LURASIDONE HYDROCHLORIDE

TABLET; ORAL

LATUDA

AB	+	SUNOVION PHARMS INC	20MG	N200603 003	Dec 07, 2011
AB	+	!	40MG	N200603 001	Oct 28, 2010
AB	+		60MG	N200603 005	Jul 12, 2013
AB	+		80MG	N200603 002	Oct 28, 2010
AB	+		120MG	N200603 004	Apr 26, 2012

PRESCRIPTION DRUG PRODUCT LIST

LURASIDONE HYDROCHLORIDE

TABLET; ORAL

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>	SUN PHARM	<u>20MG</u>	<u>A208066 001</u>	Jan 04, 2019
<u>AB</u>		<u>40MG</u>	<u>A208066 002</u>	Jan 04, 2019
<u>AB</u>		<u>60MG</u>	<u>A208066 003</u>	Jan 04, 2019
<u>AB</u>		<u>80MG</u>	<u>A208066 004</u>	Jan 04, 2019
<u>AB</u>		<u>120MG</u>	<u>A208066 005</u>	Jan 04, 2019

LUSUTROMBOPAG

TABLET; ORAL

MULPLETA

+! SHIONOGI INC 3MG N210923 001 Jul 31, 2018

LUTETIUM DOTATATE LU-177

SOLUTION; INTRAVENOUS

LUTATHERA

+! AAA USA INC 10mCi/ML N208700 001 Jan 26, 2018

MACIMORELIN ACETATE

FOR SOLUTION; ORAL

MACRILEN

+! NOVO EQ 60MG BASE/POUCH N205598 001 Dec 20, 2017

MACITENTAN

TABLET; ORAL

OPSUMIT

+! ACTELION PHARMS LTD 10MG N204410 001 Oct 18, 2013

MAFENIDE ACETATE

CREAM; TOPICAL

SULFAMYLON

+! MYLAN INSTITUTIONAL EQ 85MG BASE/GM N016763 001

FOR SOLUTION; TOPICAL

MAFENIDE ACETATE

<u>AT</u>	NOVAST LABS	<u>5%</u>	<u>A206716 001</u>	Jul 31, 2017
<u>AT</u>	PAR FORM	<u>5%</u>	<u>A201511 001</u>	Feb 12, 2013
<u>AT</u>	+! MYLAN INSTITUTIONAL	<u>5%</u>	<u>N019832 003</u>	Jun 05, 1998

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

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

PLASMA-LYTE 148 IN WATER IN PLASTIC CONTAINER

+! BAXTER HLTHCARE 30MG/100ML; 37MG/100ML; 368MG/100ML; 526MG/100ML; 502MG/100ML N017378 001

PLASMA-LYTE A IN PLASTIC CONTAINER

+! BAXTER HLTHCARE 30MG/100ML; 37MG/100ML; 368MG/100ML; 526MG/100ML; 502MG/100ML N017378 002 Nov 22, 1982

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

PRESCRIPTION DRUG PRODUCT LIST

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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 IN DEXTROSE 5% IN PLASTIC CONTAINERAP FRESENIUS KABI USA 1GM/100ML A206486 001 Mar 07, 2016AP +! HOSPIRA 1GM/100ML N020488 001 Jul 11, 1995AP HQ SPCLT PHARMA 1GM/100ML A207349 001 Mar 02, 2016AP MYLAN LABS LTD 1GM/100ML A209932 001 Sep 10, 2018MAGNESIUM SULFATE IN PLASTIC CONTAINERAP FRESENIUS KABI USA 4GM/100ML (40MG/ML) A206485 001 Mar 15, 2016AP 4GM/50ML (80MG/ML) A206485 002 Mar 15, 2016AP 2GM/50ML (40MG/ML) A206485 003 Mar 15, 2016AP 20GM/500ML (40MG/ML) A206485 004 Mar 15, 2016AP 40GM/1000ML (40MG/ML) A206485 005 Mar 15, 2016AP + HOSPIRA 2GM/50ML (40MG/ML) N020309 003 Jan 26, 2007AP +! 4GM/100ML (40MG/ML) N020309 001 Jun 24, 1994AP +! 4GM/50ML (80MG/ML) N020309 002 Jun 24, 1994AP + 20GM/500ML (40MG/ML) N020309 004 Jan 18, 1995AP + 40GM/1000ML (40MG/ML) N020309 005 Jan 18, 1995AP HQ SPCLT PHARMA 2GM/50ML (40MG/ML) A207350 001 Dec 06, 2017AP 4GM/100ML (40MG/ML) A207350 002 Dec 06, 2017AP 4GM/50ML (80MG/ML) A207350 003 Dec 06, 2017AP 20GM/500ML (40MG/ML) A207350 004 Dec 06, 2017AP 40GM/1000ML (40MG/ML) A207350 005 Dec 06, 2017AP MYLAN LABS LTD 2GM/50ML (40MG/ML) A209911 001 Sep 14, 2018AP 4GM/100ML (40MG/ML) A209911 002 Sep 14, 2018

MAGNESIUM SULFATE IN DEXTROSE 5% IN PLASTIC CONTAINER

+! HOSPIRA 2GM/100ML N020488 002 Jul 11, 1995

SOLUTION; INTRAMUSCULAR, INTRAVENOUS

MAGNESIUM SULFATEAP EXELA PHARMA SCS 5GM/10ML (500MG/ML) A206039 001 Dec 18, 2014

LLC

AP +! FRESENIUS KABI USA 5GM/10ML (500MG/ML) N019316 001 Sep 08, 1986AP +! 10GM/20ML (500MG/ML) N019316 003 Jan 29, 2016AP ! HOSPIRA 5GM/10ML (500MG/ML) A075151 001 Apr 25, 2000AP HOSPIRA INC 10GM/20ML (500MG/ML) A202411 001 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; POTASSIUM CHLORIDE; POTASSIUM PHOSPHATE, MONOBASIC; SODIUM CHLORIDE; SODIUM PHOSPHATE

SOLUTION; IRRIGATION

TIS-U-SOLAT BAXTER HLTHCARE 20MG/100ML; 40MG/100ML; 6.25MG/100ML; 800MG/100ML; 8.75MG/100ML N018508 001 Feb 19, 1982TIS-U-SOL IN PLASTIC CONTAINERAT BAXTER HLTHCARE 20MG/100ML; 40MG/100ML; 6.25MG/100ML; 800MG/100ML; 8.75MG/100ML N018336 001MAGNESIUM SULFATE; POTASSIUM SULFATE; SODIUM SULFATE

SOLUTION; ORAL

SODIUM SULFATE, POTASSIUM SULFATE AND MAGNESIUM SULFATEAA NOVEL LABS INC 1.6GM/BOT; 3.13GM/BOT; 17.5GM/BOT A202511 001 Feb 23, 2017SUPREP BOWEL PREP KITAA +! BRAINTREE LABS 1.6GM/BOT; 3.13GM/BOT; 17.5GM/BOT N022372 001 Aug 05, 2010MALATHION

LOTION; TOPICAL

MALATHION

! SUVEN LIFE 0.5% A091559 001 May 23, 2012

MANGANESE CHLORIDE

INJECTABLE; INJECTION

MANGANESE CHLORIDE IN PLASTIC CONTAINER

+! HOSPIRA EQ 0.1MG MANGANESE/ML N018962 001 Jun 26, 1986

PRESCRIPTION DRUG PRODUCT LIST

MANNITOL

INJECTABLE; INJECTION

MANNITOL 10% IN PLASTIC CONTAINER

<u>AP</u>	B BRAUN	<u>10GM/100ML</u>	<u>N020006 002</u>	Jul 26, 1993
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MANNITOL 15% IN PLASTIC CONTAINER

<u>AP</u>	B BRAUN	<u>15GM/100ML</u>	<u>N020006 003</u>	Jul 26, 1993
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MANNITOL 20% IN PLASTIC CONTAINER

<u>AP</u>	B BRAUN	<u>20GM/100ML</u>	<u>N020006 004</u>	Jul 26, 1993
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<u>AP</u>	ICU MEDICAL INC	<u>20GM/100ML</u>	<u>N019603 004</u>	Jan 08, 1990
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MANNITOL 25%

<u>AP</u>	FRESENIUS KABI USA	<u>12.5GM/50ML</u>	<u>A080677 001</u>	
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<u>AP</u>	HOSPIRA	<u>12.5GM/50ML</u>	<u>N016269 006</u>	Aug 25, 1994
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<u>AP</u>	INTL MEDICATION	<u>12.5GM/50ML</u>	<u>A083051 001</u>	
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MANNITOL 5% IN PLASTIC CONTAINER

<u>AP</u>	B BRAUN	<u>5GM/100ML</u>	<u>N020006 001</u>	Jul 26, 1993
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OSMITROL 10% IN WATER

<u>AP</u>	BAXTER HLTHCARE	<u>10GM/100ML</u>	<u>N013684 002</u>	
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OSMITROL 10% IN WATER IN PLASTIC CONTAINER

<u>AP</u>	BAXTER HLTHCARE	<u>10GM/100ML</u>	<u>N013684 006</u>	
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OSMITROL 15% IN WATER

<u>AP</u>	BAXTER HLTHCARE	<u>15GM/100ML</u>	<u>N013684 004</u>	
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OSMITROL 15% IN WATER IN PLASTIC CONTAINER

<u>AP</u>	BAXTER HLTHCARE	<u>15GM/100ML</u>	<u>N013684 008</u>	
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OSMITROL 20% IN WATER

<u>AP</u>	BAXTER HLTHCARE	<u>20GM/100ML</u>	<u>N013684 003</u>	
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OSMITROL 20% IN WATER IN PLASTIC CONTAINER

<u>AP</u>	BAXTER HLTHCARE	<u>20GM/100ML</u>	<u>N013684 007</u>	
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OSMITROL 5% IN WATER

<u>AP</u>	BAXTER HLTHCARE	<u>5GM/100ML</u>	<u>N013684 001</u>	
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OSMITROL 5% IN WATER IN PLASTIC CONTAINER

<u>AP</u>	BAXTER HLTHCARE	<u>5GM/100ML</u>	<u>N013684 005</u>	
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POWDER; INHALATION

ARIDOL KIT

+	PHARMAXIS LTD	N/A, 5MG, 10MG, 20MG, 40MG	N022368 001	Oct 05, 2010
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SOLUTION; IRRIGATION

RESECTISOL IN PLASTIC CONTAINER

	B BRAUN	5GM/100ML	N016772 002	
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MANNITOL; SORBITOL

SOLUTION; IRRIGATION

SORBITOL-MANNITOL IN PLASTIC CONTAINER

+	ICU MEDICAL INC	540MG/100ML; 2.7GM/100ML	N018316 001	
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MAPROTILINE HYDROCHLORIDE

TABLET; ORAL

MAPROTILINE HYDROCHLORIDE

	MYLAN	25MG	A072285 002	Oct 03, 1988
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!		50MG	A072285 001	Oct 03, 1988
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		75MG	A072285 003	Oct 03, 1988
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MARAVIROC

SOLUTION; ORAL

SELZENTRY

+	VIIV HLTHCARE	20MG/ML	N208984 001	Nov 04, 2016
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TABLET; ORAL

SELZENTRY

+	VIIV HLTHCARE	25MG	N022128 003	Nov 04, 2016
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+		75MG	N022128 004	Nov 04, 2016
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+		150MG	N022128 001	Aug 06, 2007
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+	!	300MG	N022128 002	Aug 06, 2007
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MEBENDAZOLE

TABLET, CHEWABLE; ORAL

EMVERM

!	IMPAX LABS INC	100MG	A073580 001	Jan 04, 1995
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MECAMYLAMINE HYDROCHLORIDE

TABLET; ORAL

MECAMYLAMINE HYDROCHLORIDE

!	NEXGEN PHARMA	2.5MG	A204054 001	Mar 19, 2013
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PRESCRIPTION DRUG PRODUCT LIST

MECASERMIN RECOMBINANT

INJECTABLE; SUBCUTANEOUS

INCRELEX

+! IPSEN INC 40MG/4ML (10MG/ML) N021839 001 Aug 30, 2005

MECHLORETHAMINE HYDROCHLORIDE

GEL; TOPICAL

VALCHLOR

+! HELSINN EQ 0.016% BASE N202317 001 Aug 23, 2013

MECLIZINE HYDROCHLORIDE

TABLET; ORAL

ANTIVERT

AA + CASPER PHARMA LLC 12.5MG **N010721 006**
AA + 25MG **N010721 004**
AA + 50MG **N010721 001** Jan 20, 1982

MECLIZINE HYDROCHLORIDE

AA ! AMNEAL PHARMS 12.5MG **A201451 001** Feb 23, 2011
AA ! 25MG **A201451 002** Feb 23, 2011
AA AUREX 12.5MG **A087127 001**
AA 25MG **A087128 001**
AA EPIC PHARMA LLC 12.5MG **A200294 001** Apr 13, 2012
AA 25MG **A200294 002** Apr 13, 2012
AA INDICUS PHARMA 12.5MG **A205136 001** Feb 22, 2019
AA 25MG **A205136 002** Feb 22, 2019
AA INVATECH 25MG **A084092 003** May 22, 1989
AA JUBILANT CADISTA 12.5MG **A040659 001** Jun 04, 2010
AA 25MG **A040659 002** Jun 04, 2010
AA SANDOZ 12.5MG **A084843 002** May 22, 1989
 INDICUS PHARMA 50MG **A205136 003** Feb 22, 2019

TABLET, CHEWABLE; ORAL

ANTIVERT

AA + 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

AB +! PHARMACIA AND 150MG/ML **N020246 001** Oct 29, 1992
 UPJOHN

MEDROXYPROGESTERONE ACETATE

AB AMPHASTAR PHARMS 150MG/ML **A077235 001** Nov 28, 2017
 INC
AB 150MG/ML **A077334 001** Nov 28, 2017
AB MYLAN LABS LTD 150MG/ML **A210227 001** Oct 12, 2018
AB SUN PHARM 150MG/ML **A210760 001** May 01, 2019
AB 150MG/ML **A210761 001** Apr 24, 2019
AB TEVA PHARMS USA 150MG/ML **A076553 001** Jul 28, 2004

DEPO-PROVERA

+! PHARMACIA AND 400MG/ML N012541 003
 UPJOHN

INJECTABLE; SUBCUTANEOUS

DEPO-SUBQ PROVERA 104

+! PHARMACIA AND 104MG/0.65ML N021583 001 Dec 17, 2004
 UPJOHN

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 + PHARMACIA AND 2.5MG **N011839 001**
 UPJOHN
AB + 5MG **N011839 003**
AB +! 10MG **N011839 004**

PRESCRIPTION DRUG PRODUCT LIST

3-279 (of 453)

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	+ !	SHIONOGI INC	250MG	N015034	003
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MEFLOQUINE HYDROCHLORIDE

TABLET; ORAL

MEFLOQUINE HYDROCHLORIDE

!	BARR	250MG	A076392	001	Dec 29, 2003
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MEGESTROL ACETATE

SUSPENSION; ORAL

MEGACE ES

AB	+ !	ENDO PHARMS INC	125MG/ML	N021778	001	Jul 05, 2005
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MEGESTROL ACETATE

AB		BRECKENRIDGE	125MG/ML	A204688	001	Dec 01, 2017
AB		HI TECH	40MG/ML	A203960	001	Jun 09, 2017
AB	!	PAR PHARM	40MG/ML	A075671	001	Jul 25, 2001
AB		TEVA PHARMS	40MG/ML	A075681	001	May 05, 2003
AB		TWI PHARMS	125MG/ML	A203139	001	Aug 27, 2014
AB		WOCKHARDT BIO AG	40MG/ML	A076721	001	Nov 01, 2004

TABLET; ORAL

MEGESTROL ACETATE

AB		BARR	20MG	A074621	002	Aug 16, 1996
AB			40MG	A074621	001	Nov 30, 1995
AB		HIKMA	20MG	A074458	001	Sep 29, 1995
AB			40MG	A074458	002	Sep 29, 1995
AB		PAR PHARM	20MG	A072422	001	Aug 08, 1988
AB	!		40MG	A072423	001	Aug 08, 1988

MELOXICAM

CAPSULE; ORAL

VIVLODEX

+ ZYLA

+!

N207233 001 Oct 22, 2015

N207233 002 Oct 22, 2015

SOLUTION; INTRAVENOUS

ANJESO

+! BAUDAX

30MG/ML (30MG/ML)

N210583 001 Feb 20, 2020

TABLET; ORAL

MELOXICAM

AB		APOTEX INC	7.5MG	A077882	001	Jul 20, 2006
AB			15MG	A077882	002	Jul 20, 2006
AB		AUROBINDO PHARMA	7.5MG	A078008	001	Oct 02, 2006
AB			15MG	A078008	002	Oct 02, 2006
AB		BRECKENRIDGE PHARM	7.5MG	A077920	001	Jul 19, 2006
AB			15MG	A077920	002	Jul 19, 2006
AB		CIPLA	7.5MG	A077929	001	Jul 19, 2006
AB			15MG	A077929	002	Jul 19, 2006
AB		DR REDDYS LABS INC	7.5MG	A077931	001	Jul 25, 2006
AB			15MG	A077931	002	Jul 25, 2006
AB		GLENMARK GENERICS	7.5MG	A077932	001	Jul 19, 2006
AB			15MG	A077932	002	Jul 19, 2006
AB		LUPIN PHARMS	7.5MG	A077944	001	Jul 19, 2006
AB			15MG	A077944	002	Jul 19, 2006
AB		PURACAP PHARM	7.5MG	A077938	001	Jul 19, 2006
AB			15MG	A077938	002	Jul 19, 2006
AB		STRIDES PHARMA	7.5MG	A077928	001	May 13, 2009
AB			15MG	A077928	002	May 13, 2009
AB		TARO	7.5MG	A078102	001	Nov 07, 2006
AB			15MG	A078102	002	Nov 07, 2006
AB		UNICHEM	7.5MG	A077927	001	Dec 20, 2006
AB			15MG	A077927	002	Dec 20, 2006
AB		YUNG SHIN PHARM	7.5MG	A077918	001	Dec 07, 2006
AB			15MG	A077918	002	Dec 07, 2006
AB		ZYDUS PHARMS USA	7.5MG	A077921	001	Jul 19, 2006
AB			15MG	A077921	002	Jul 19, 2006

MOBIC

AB	+	BOEHRINGER INGELHEIM	7.5MG	N020938	001	Apr 13, 2000
AB	+ !		15MG	N020938	002	Aug 23, 2000

PRESCRIPTION DRUG PRODUCT LIST

MELOXICAM

TABLET, ORALLY DISINTEGRATING;ORAL
 QMIIZ ODT
 + TERSERA THERAPS LLC 7.5MG
 +! 15MG

N211210 001 Oct 19, 2018
 N211210 002 Oct 19, 2018

MELPHALAN

TABLET;ORAL

ALKERAN

AB +! APOTEX **2MG**

N014691 002

MELPHALAN

AB ALVOGEN **2MG**

A207809 001 Mar 22, 2017

MELPHALAN HYDROCHLORIDE

INJECTABLE; INJECTION

ALKERAN

AP + APOTEX **EQ 50MG BASE/VIAL**

N020207 001 Nov 18, 1992

MELPHALAN HYDROCHLORIDE

AP ACTAVIS LLC **EQ 50MG BASE/VIAL**

A206018 001 Dec 19, 2016

AP ALVOGEN INC **EQ 50MG BASE/VIAL**

A204817 001 May 17, 2019

AP DR REDDYS LABS LTD **EQ 50MG BASE/VIAL**

A203655 001 Dec 08, 2017

AP FRESENIUS KABI USA **EQ 50MG BASE/VIAL**

A203393 001 Dec 22, 2017

AP GLAND PHARMA LTD **EQ 50MG BASE/VIAL**

A209826 001 May 28, 2019

AP INGENUS PHARMS LLC **EQ 50MG BASE/VIAL**

A210947 001 Feb 18, 2020

AP ! MYLAN INSTITUTIONAL **EQ 50MG BASE/VIAL**

A090270 001 Jun 09, 2009

AP SAGENT PHARMS INC **EQ 50MG BASE/VIAL**

A201379 001 Feb 28, 2017

AP WEST-WARD PHARMS **EQ 50MG BASE/VIAL**

A090303 001 Oct 28, 2010

POWDER; INTRAVENOUS

EVOMELA

AP +! ACROTECH **EQ 50MG BASE/VIAL**

N207155 001 Mar 10, 2016

MELPHALAN HYDROCHLORIDE

AP ACTAVIS LLC **EQ 50MG BASE/VIAL**

A209323 001 Mar 06, 2020

MEMANTINE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE;ORAL

MEMANTINE HYDROCHLORIDE

AB AMNEAL PHARMS **7MG**

A205825 001 Oct 12, 2016

AB **14MG**

A205825 002 Oct 12, 2016

AB **21MG**

A205825 003 Oct 12, 2016

AB **28MG**

A205825 004 Oct 12, 2016

AB ANCHEN PHARMS **7MG**

A205784 001 Jun 09, 2017

AB **14MG**

A205784 002 Jun 09, 2017

AB **21MG**

A205784 003 Jun 09, 2017

AB **28MG**

A205784 004 Jun 09, 2017

AB ANI PHARMS INC **7MG**

A205365 001 Feb 28, 2020

AB **14MG**

A205365 002 Feb 28, 2020

AB **21MG**

A205365 003 Feb 28, 2020

AB **28MG**

A205365 004 Feb 28, 2020

AB APOTEX **7MG**

A206135 001 Nov 22, 2016

AB **14MG**

A206135 002 Nov 22, 2016

AB **21MG**

A206135 003 Nov 22, 2016

AB **28MG**

A206135 004 Nov 22, 2016

AB LUPIN LTD **7MG**

A206028 001 Sep 28, 2016

AB **14MG**

A206028 002 Sep 28, 2016

AB **21MG**

A206028 003 Sep 28, 2016

AB **28MG**

A206028 004 Sep 28, 2016

AB MYLAN **7MG**

A206032 001 Sep 28, 2016

AB **14MG**

A206032 002 Sep 28, 2016

AB **21MG**

A206032 003 Sep 28, 2016

AB **28MG**

A206032 004 Sep 28, 2016

AB SUN PHARM **7MG**

A205905 001 Sep 28, 2016

AB **14MG**

A205905 002 Sep 28, 2016

AB **21MG**

A205905 003 Sep 28, 2016

AB **28MG**

A205905 004 Sep 28, 2016

AB ZYDUS PHARMS **7MG**

A203293 001 Aug 03, 2017

AB **14MG**

A203293 002 Aug 03, 2017

AB **21MG**

A203293 003 Aug 03, 2017

AB **28MG**

A203293 004 Aug 03, 2017

NAMENDA XR

AB + FOREST LABS LLC **7MG**

N022525 001 Jun 21, 2010

AB + **14MG**

N022525 002 Jun 21, 2010

AB + **21MG**

N022525 003 Jun 21, 2010

AB +! **28MG**

N022525 004 Jun 21, 2010

PRESCRIPTION DRUG PRODUCT LIST

MEMANTINE HYDROCHLORIDE

SOLUTION;ORAL

MEMANTINE HYDROCHLORIDE

<u>AA</u>	APOTEX	<u>2MG/ML</u>	<u>A209955</u>	<u>001</u>	Feb 09, 2018
<u>AA</u>	! LANNETT CO INC	<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

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 PHARMS LTD	<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>	CADILA	<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
<u>AB</u>	CELLTRION	<u>5MG</u>	<u>A090073</u>	<u>001</u>	Sep 04, 2015
<u>AB</u>		<u>10MG</u>	<u>A090073</u>	<u>002</u>	Sep 04, 2015
<u>AB</u>	CSEPC OUYI	<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>	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>	HIKMA PHARMS	<u>5MG</u>	<u>A208173</u>	<u>001</u>	Feb 28, 2020
<u>AB</u>		<u>10MG</u>	<u>A208173</u>	<u>002</u>	Feb 28, 2020
<u>AB</u>	JUBILANT GENERICS	<u>5MG</u>	<u>A091585</u>	<u>001</u>	Oct 13, 2015
<u>AB</u>		<u>10MG</u>	<u>A091585</u>	<u>002</u>	Oct 13, 2015
<u>AB</u>	LANNETT CO INC	<u>5MG</u>	<u>A207236</u>	<u>001</u>	Nov 10, 2016
<u>AB</u>		<u>10MG</u>	<u>A207236</u>	<u>002</u>	Nov 10, 2016
<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>	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>	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>	TEVA PHARMS	<u>5MG</u>	<u>A090052</u>	<u>001</u>	Oct 25, 2011
<u>AB</u>		<u>10MG</u>	<u>A090052</u>	<u>002</u>	Oct 25, 2011
<u>AB</u>	TORRENT	<u>5MG</u>	<u>A200155</u>	<u>001</u>	Oct 13, 2015
<u>AB</u>		<u>10MG</u>	<u>A200155</u>	<u>002</u>	Oct 13, 2015
<u>AB</u>	UNICHEM LABS LTD	<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

NAMENDA

<u>AB</u>	+ ALLERGAN	<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

MENOTROPINS (FSH;LH)

INJECTABLE;SUBCUTANEOUS

MENOPUR

+!	FERRING	75 IU/VIAL;75 IU/VIAL	N021663	001	Oct 29, 2004
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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>A080445</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>A080445</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>A080445</u>	<u>009</u>	
<u>AP</u>		<u>100MG/ML</u>	<u>A080445</u>	<u>004</u>	

PRESCRIPTION DRUG PRODUCT LIST

MEPERIDINE HYDROCHLORIDE

INJECTABLE; INJECTION

MEPERIDINE HYDROCHLORIDE

<u>AP</u>		<u>100MG/ML</u>	<u>A080455</u>	<u>010</u>	
	MEPERIDINE HYDROCHLORIDE PRESERVATIVE FREE				
	! WEST-WARD PHARMS	10MG/ML	A081002	001	Jul 30, 1993
	INT				
	SYRUP; ORAL				
	MEPERIDINE HYDROCHLORIDE				
	! HIKMA	50MG/5ML	A088744	001	Jan 30, 1985
	TABLET; ORAL				
	<u>MEPERIDINE HYDROCHLORIDE</u>				
<u>AA</u>	! EPIC PHARMA	<u>50MG</u>	<u>A040331</u>	<u>001</u>	May 28, 1999
<u>AA</u>	!	<u>100MG</u>	<u>A040331</u>	<u>002</u>	May 28, 1999
<u>AA</u>	VINTAGE PHARMS	<u>50MG</u>	<u>A040191</u>	<u>001</u>	Dec 17, 1998
<u>AA</u>		<u>100MG</u>	<u>A040191</u>	<u>002</u>	Dec 17, 1998

MEPIVACAINE HYDROCHLORIDE

INJECTABLE; INJECTION

CARBOCAINE

<u>AP</u>	+! HOSPIRA	<u>1%</u>	<u>N012250</u>	<u>001</u>	
<u>AP</u>	+!	<u>1.5%</u>	<u>N012250</u>	<u>005</u>	
<u>AP</u>	+!	<u>2%</u>	<u>N012250</u>	<u>002</u>	
	<u>POLOCAINE</u>				
<u>AP</u>	FRESENIUS KABI USA	<u>1%</u>	<u>A089407</u>	<u>001</u>	Dec 01, 1986
<u>AP</u>		<u>2%</u>	<u>A089410</u>	<u>001</u>	Dec 01, 1986
	<u>POLOCAINE-MPF</u>				
<u>AP</u>	FRESENIUS KABI USA	<u>1%</u>	<u>A089406</u>	<u>001</u>	Dec 01, 1986
<u>AP</u>		<u>1.5%</u>	<u>A089408</u>	<u>001</u>	Dec 01, 1986
<u>AP</u>		<u>2%</u>	<u>A089409</u>	<u>001</u>	Dec 01, 1986
	<u>SCANDONEST PLAIN</u>				
<u>AP</u>	! DEPROCO	<u>3%</u>	<u>A088387</u>	<u>001</u>	Oct 10, 1984

MEPROBAMATE

TABLET; ORAL

MEPROBAMATE

<u>AA</u>	ALEMBIC PHARMS LTD	<u>200MG</u>	<u>A090122</u>	<u>001</u>	Feb 18, 2009
<u>AA</u>		<u>400MG</u>	<u>A090122</u>	<u>002</u>	Feb 18, 2009
<u>AA</u>	INVAGEN PHARMS	<u>200MG</u>	<u>A040797</u>	<u>001</u>	Feb 27, 2008
<u>AA</u>		<u>400MG</u>	<u>A040797</u>	<u>002</u>	Feb 27, 2008
<u>AA</u>	! WATSON LABS	<u>200MG</u>	<u>A083304</u>	<u>001</u>	
<u>AA</u>	!	<u>400MG</u>	<u>A083308</u>	<u>001</u>	

MERCAPTOPYRINE

SUSPENSION; ORAL

PURIXAN

+! NOVA LABS LTD 20MG/ML

N205919 001 Apr 28, 2014

TABLET; ORAL

MERCAPTOPYRINE

<u>AB</u>	DR REDDYS LABS SA	<u>50MG</u>	<u>A040461</u>	<u>001</u>	Feb 11, 2004
<u>AB</u>	! HIKMA	<u>50MG</u>	<u>A040528</u>	<u>001</u>	Feb 13, 2004
<u>AB</u>	MYLAN	<u>50MG</u>	<u>A040594</u>	<u>001</u>	Jul 01, 2005
	<u>PURINETHOL</u>				
<u>AB</u>	+ STASON PHARMS	<u>50MG</u>	<u>N009053</u>	<u>002</u>	

MEROPENEM

INJECTABLE; INJECTION

MEROPENEM

<u>AP</u>	ACS DOBFAR	<u>500MG/VIAL</u>	<u>A091404</u>	<u>001</u>	Oct 26, 2011
<u>AP</u>		<u>1GM/VIAL</u>	<u>A091404</u>	<u>002</u>	Oct 26, 2011
<u>AP</u>	ACS DOBFAR SPA	<u>500MG/VIAL</u>	<u>A204139</u>	<u>001</u>	Jun 09, 2016
<u>AP</u>		<u>1GM/VIAL</u>	<u>A204139</u>	<u>002</u>	Jun 09, 2016
<u>AP</u>	AMNEAL PHARMS	<u>500MG/VIAL</u>	<u>A205883</u>	<u>001</u>	Apr 12, 2016
<u>AP</u>		<u>1GM/VIAL</u>	<u>A205883</u>	<u>002</u>	Apr 12, 2016
<u>AP</u>	AUROBINDO PHARMA LTD	<u>500MG/VIAL</u>	<u>A205835</u>	<u>001</u>	Mar 27, 2017
<u>AP</u>		<u>1GM/VIAL</u>	<u>A205835</u>	<u>002</u>	Mar 27, 2017
<u>AP</u>	DAEWOONG PHARM CO	<u>500MG/VIAL</u>	<u>A204854</u>	<u>001</u>	Dec 18, 2015
<u>AP</u>		<u>1GM/VIAL</u>	<u>A204854</u>	<u>002</u>	Dec 18, 2015
<u>AP</u>	GLAND PHARMA LTD	<u>500MG/VIAL</u>	<u>A206141</u>	<u>001</u>	Jun 08, 2016
<u>AP</u>		<u>1GM/VIAL</u>	<u>A206141</u>	<u>002</u>	Jun 08, 2016
<u>AP</u>	HQ SPCLT PHARMA	<u>500MG/VIAL</u>	<u>A210773</u>	<u>001</u>	Aug 16, 2019
<u>AP</u>		<u>1GM/VIAL</u>	<u>A210773</u>	<u>002</u>	Aug 16, 2019
<u>AP</u>	SAVIOR LIFETEC CORP	<u>500MG/VIAL</u>	<u>A206086</u>	<u>001</u>	Apr 19, 2016
<u>AP</u>		<u>1GM/VIAL</u>	<u>A206086</u>	<u>002</u>	Apr 19, 2016

PRESCRIPTION DRUG PRODUCT LIST

MEROPENEM

INJECTABLE; INJECTION

MERREM

AP	+	PFIZER	500MG/VIAL	N050706	003	Jun 21, 1996
AP	+		1GM/VIAL	N050706	001	Jun 21, 1996

POWDER; INTRAVENOUS

MEROPENEM AND SODIUM CHLORIDE IN DUPLEX CONTAINER

B BRAUN MEDICAL INC	500MG/VIAL	N202106	001	Apr 30, 2015
	1GM/VIAL	N202106	002	Apr 30, 2015

MEROPENEM; VABORBACTAM

POWDER; INTRAVENOUS

VABOMERE

+	REMPEX PHARMS	1GM/VIAL;1GM/VIAL	N209776	001	Aug 29, 2017
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MESALAMINE

CAPSULE, DELAYED RELEASE; ORAL

DELZICOL

AB	+	APIL	400MG	N204412	001	Feb 01, 2013
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MESALAMINE

AB		TEVA PHARMS USA	400MG	A207873	001	May 09, 2019
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CAPSULE, EXTENDED RELEASE; ORAL

APRISO

AB	+	VALEANT PHARMS INTL	375MG	N022301	001	Oct 31, 2008
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MESALAMINE

AB		MYLAN	375MG	A207271	001	Nov 20, 2019
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PENTASA

+	SHIRE	250MG	N020049	001	May 10, 1993
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+		500MG	N020049	002	Jul 08, 2004
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ENEMA; RECTAL

MESALAMINE

AB		PERRIGO ISRAEL	4GM/60ML	A076751	001	Sep 17, 2004
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ROWASA

AB	+	MYLAN SPECIALITY LP	4GM/60ML	N019618	001	Dec 24, 1987
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SFROWASA

AB	+	MYLAN SPECIALITY LP	4GM/60ML	N019618	002	Jun 20, 2008
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SUPPOSITORY; RECTAL

CANASA

AB	+	ALLERGAN	1GM	N021252	002	Nov 05, 2004
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MESALAMINE

AB		AMNEAL PHARMS LLC	1GM	A210509	001	Jan 02, 2020
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AB		AMRING PHARMS	1GM	A208362	001	Jun 21, 2019
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AB		MYLAN	1GM	A204354	001	Nov 24, 2015
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AB		PHARM SOURCING	1GM	A207448	001	Apr 19, 2019
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AB		SANDOZ INC	1GM	A202065	001	Jun 12, 2019
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AB		ZYDUS PHARMS	1GM	A208953	001	Feb 12, 2020
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TABLET, DELAYED RELEASE; ORAL

ASACOL HD

AB	+	APIL	800MG	N021830	001	May 29, 2008
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LIALDA

AB	+	SHIRE	1.2GM	N022000	001	Jan 16, 2007
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MESALAMINE

AB		ACTAVIS LABS FL	1.2GM	A203817	001	Mar 23, 2018
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AB		MYLAN	1.2GM	A203574	001	Nov 20, 2018
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AB		SUN PHARM	1.2GM	A211858	001	Jan 25, 2019
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AB		ZYDUS PHARMS	800MG	A203286	001	Jul 21, 2017
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AB			1.2GM	A091640	001	Jun 05, 2017
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MESNA

INJECTABLE; INTRAVENOUS

MESNA

AP		FRESENIUS KABI USA	100MG/ML	A075811	001	Apr 26, 2001
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AP		GLAND PHARMA LTD	100MG/ML	A206992	001	Dec 18, 2017
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AP		SAGENT PHARMS INC	100MG/ML	A090913	001	Apr 13, 2010
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AP		TEVA PHARMS USA	100MG/ML	A075764	001	Apr 27, 2001
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AP		WEST-WARD PHARMS	100MG/ML	A075739	001	Jan 09, 2004
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MESNEX

AP	+	BAXTER HLTHCARE	100MG/ML	N019884	001	Dec 30, 1988
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TABLET; ORAL

MESNEX

+	BAXTER HLTHCARE	400MG	N020855	001	Mar 21, 2002
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PRESCRIPTION DRUG PRODUCT LIST

METAPROTERENOL SULFATE

SYRUP;ORAL

METAPROTERENOL SULFATE

! LANNETT CO INC 10MG/5ML

A073632 001 Jul 22, 1992

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>	APPCO	<u>800MG</u>	<u>A208774</u>	<u>001</u>	Sep 24, 2018
<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

SKELAXIN

<u>AB</u>	+! KING PHARMS	<u>800MG</u>	<u>N013217</u>	<u>003</u>	Aug 30, 2002
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METFORMIN HYDROCHLORIDE

FOR SUSPENSION, EXTENDED RELEASE;ORAL

RIOMET ER

+! SUN PHARM 500MG/5ML

N212595 001 Aug 29, 2019

SOLUTION;ORAL

METFORMIN HYDROCHLORIDE

<u>AB</u>	SAPTALIS PHARMS	<u>500MG/5ML</u>	<u>A211309</u>	<u>001</u>	Mar 03, 2020
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RIOMET

<u>AB</u>	+! RANBAXY	<u>500MG/5ML</u>	<u>N021591</u>	<u>001</u>	Sep 11, 2003
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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>	AMNEAL PHARMS NY	<u>500MG</u>	<u>A077880</u>	<u>001</u>	Jun 05, 2006
<u>AB</u>		<u>850MG</u>	<u>A077880</u>	<u>002</u>	Jun 05, 2006
<u>AB</u>		<u>1GM</u>	<u>A077880</u>	<u>003</u>	Jun 05, 2006
<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 LIFE SCI	<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>	GLENMARK GENERICS	<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>	HERITAGE PHARMA	<u>1GM</u>	<u>A075978</u>	<u>003</u>	Nov 05, 2002
<u>AB</u>		<u>500MG</u>	<u>A075978</u>	<u>001</u>	Jan 25, 2002
<u>AB</u>		<u>850MG</u>	<u>A075978</u>	<u>002</u>	Jan 25, 2002
<u>AB</u>	INDICUS PHARMA	<u>500MG</u>	<u>A079148</u>	<u>001</u>	Nov 25, 2008
<u>AB</u>		<u>850MG</u>	<u>A079148</u>	<u>002</u>	Nov 25, 2008
<u>AB</u>		<u>1GM</u>	<u>A079148</u>	<u>003</u>	Nov 25, 2008
<u>AB</u>	LAURUS LABS LTD	<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>	MACLEODS PHARMS LTD	<u>500MG</u>	<u>A205330</u>	<u>001</u>	Oct 31, 2017

PRESCRIPTION DRUG PRODUCT LIST

METFORMIN HYDROCHLORIDE

TABLET; ORAL

METFORMIN HYDROCHLORIDE

<u>AB</u>		<u>850MG</u>	<u>A205330 002</u>	Oct 31, 2017
<u>AB</u>		<u>1GM</u>	<u>A205330 003</u>	Oct 31, 2017
<u>AB</u>	MARKSANS PHARMA	<u>500MG</u>	<u>A090888 001</u>	Mar 12, 2012
<u>AB</u>		<u>850MG</u>	<u>A090888 002</u>	Mar 12, 2012
<u>AB</u>		<u>1GM</u>	<u>A090888 003</u>	Mar 12, 2012
<u>AB</u>	MYLAN	<u>500MG</u>	<u>A075973 001</u>	Jan 25, 2002
<u>AB</u>		<u>850MG</u>	<u>A075973 002</u>	Jan 25, 2002
<u>AB</u>		<u>1GM</u>	<u>A075973 003</u>	Jan 25, 2002
<u>AB</u>	SANDOZ	<u>500MG</u>	<u>A075965 001</u>	Jan 25, 2002
<u>AB</u>		<u>850MG</u>	<u>A075965 002</u>	Jan 25, 2002
<u>AB</u>		<u>1GM</u>	<u>A075965 003</u>	Jan 25, 2002
<u>AB</u>	SCIEGEN PHARMS INC	<u>500MG</u>	<u>A203769 001</u>	Sep 11, 2013
<u>AB</u>		<u>850MG</u>	<u>A203769 002</u>	Sep 11, 2013
<u>AB</u>		<u>1GM</u>	<u>A203769 003</u>	Sep 11, 2013
<u>AB</u>	SUN PHARM INDS INC	<u>500MG</u>	<u>A075967 001</u>	Jan 29, 2002
<u>AB</u>		<u>850MG</u>	<u>A075967 002</u>	Jan 29, 2002
<u>AB</u>		<u>1GM</u>	<u>A075967 003</u>	Jan 29, 2002
<u>AB</u>	TORRENT PHARMS	<u>500MG</u>	<u>A077711 001</u>	Jan 24, 2007
<u>AB</u>		<u>850MG</u>	<u>A077711 002</u>	Jan 24, 2007
<u>AB</u>		<u>1GM</u>	<u>A077711 003</u>	Jan 24, 2007
<u>AB</u>	ZYDUS HLTHCARE	<u>500MG</u>	<u>A203686 001</u>	Aug 28, 2014
<u>AB</u>		<u>850MG</u>	<u>A203686 002</u>	Aug 28, 2014
<u>AB</u>		<u>1GM</u>	<u>A203686 003</u>	Aug 28, 2014
<u>AB</u>	ZYDUS PHARMS USA	<u>500MG</u>	<u>A077064 001</u>	Apr 18, 2005
<u>AB</u>		<u>850MG</u>	<u>A077064 002</u>	Apr 18, 2005
<u>AB</u>		<u>1GM</u>	<u>A077064 003</u>	Apr 18, 2005
	CHARTWELL LIFE SCI	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 001</u>	Apr 12, 2005
<u>AB</u>	ALKEM LABS LTD	<u>750MG</u>	<u>A206145 002</u>	Oct 22, 2018
<u>AB</u>	AMNEAL PHARMS NY	<u>750MG</u>	<u>A078596 002</u>	Jan 03, 2008
<u>AB</u>	APOTEX	<u>750MG</u>	<u>A076706 002</u>	Dec 29, 2005
<u>AB</u>	AUROBINDO PHARMA LTD	<u>750MG</u>	<u>A079118 002</u>	Jul 20, 2012
<u>AB</u>	BARR	<u>750MG</u>	<u>A076863 001</u>	Oct 14, 2004
<u>AB</u>	BEXIMCO PHARMS USA	<u>750MG</u>	<u>A207427 002</u>	Dec 13, 2016
<u>AB</u>	CADILA	<u>750MG</u>	<u>A077078 001</u>	Apr 21, 2005
<u>AB</u>	CSPEC OUYI	<u>750MG</u>	<u>A078321 002</u>	Apr 17, 2008
<u>AB</u>	GRANULES INDIA LTD	<u>750MG</u>	<u>A209313 002</u>	Mar 16, 2018
<u>AB</u>	INTELLIPHARMACEUTICS	<u>750MG</u>	<u>A202306 002</u>	Feb 23, 2017
<u>AB</u>	MACLEODS PHARMS LTD	<u>750MG</u>	<u>A206955 002</u>	Dec 07, 2016
<u>AB</u>	MARKSANS PHARMA	<u>750MG</u>	<u>A090295 002</u>	Apr 29, 2016
<u>AB</u>	NOSTRUM PHARMS LLC	<u>750MG</u>	<u>A076756 002</u>	Dec 12, 2011
<u>AB</u>	PRINSTON INC	<u>750MG</u>	<u>A208880 002</u>	Sep 10, 2018
<u>AB</u>	SUN PHARM INDS (IN)	<u>750MG</u>	<u>A077336 002</u>	Feb 09, 2006
<u>AB</u>	TEVA	<u>750MG</u>	<u>A076864 001</u>	Apr 12, 2005
<u>AB1</u>	ACTAVIS LABS FL INC	<u>500MG</u>	<u>A076172 001</u>	Jun 16, 2004
<u>AB1</u>	ALIGNSCIENCE PHARMA	<u>500MG</u>	<u>A209303 001</u>	Mar 19, 2018
<u>AB1</u>	ALKEM LABS LTD	<u>500MG</u>	<u>A206145 001</u>	Oct 22, 2018
<u>AB1</u>	AMNEAL PHARMS NY	<u>500MG</u>	<u>A078596 001</u>	Jan 03, 2008
<u>AB1</u>	APOTEX	<u>500MG</u>	<u>A076706 001</u>	Dec 14, 2004
<u>AB1</u>	AUROBINDO PHARMA LTD	<u>500MG</u>	<u>A079118 001</u>	Jul 20, 2012
<u>AB1</u>	BEXIMCO PHARMS USA	<u>500MG</u>	<u>A207427 001</u>	Dec 13, 2016
<u>AB1</u>	CADILA	<u>500MG</u>	<u>A077060 001</u>	Apr 20, 2005
<u>AB1</u>	CSPEC OUYI	<u>500MG</u>	<u>A078321 001</u>	Apr 17, 2008
<u>AB1</u>	GRANULES INDIA LTD	<u>500MG</u>	<u>A209313 001</u>	Mar 16, 2018
<u>AB1</u>	INTELLIPHARMACEUTICS	<u>500MG</u>	<u>A202306 001</u>	Feb 23, 2017
<u>AB1</u>	INVENTIA	<u>500MG</u>	<u>A201991 001</u>	Jan 18, 2012
<u>AB1</u>	MACLEODS PHARMS LTD	<u>500MG</u>	<u>A206955 001</u>	Dec 07, 2016
<u>AB1</u>	MARKSANS PHARMA	<u>500MG</u>	<u>A090295 001</u>	Apr 29, 2016
<u>AB1</u>	NOSTRUM PHARMS LLC	<u>500MG</u>	<u>A076756 001</u>	Jul 26, 2006
<u>AB1</u>	PRINSTON INC	<u>500MG</u>	<u>A208880 001</u>	Sep 10, 2018
<u>AB1</u>	SANDOZ	<u>500MG</u>	<u>A076873 001</u>	Dec 14, 2004
<u>AB1</u>	SUN PHARM INDS (IN)	<u>500MG</u>	<u>A077336 001</u>	Feb 09, 2006
<u>AB1</u>	TEVA	<u>500MG</u>	<u>A076269 001</u>	Jun 18, 2004

PRESCRIPTION DRUG PRODUCT LIST

METFORMIN HYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL

METFORMIN HYDROCHLORIDE

<u>AB1</u>	TORRENT	<u>500MG</u>	<u>A090014 001</u>	Dec 30, 2009
<u>AB2</u>	LUPIN LTD	<u>500MG</u>	<u>A090692 001</u>	Jun 29, 2011
<u>AB2</u>	!	<u>1GM</u>	<u>A090692 002</u>	Jun 29, 2011
<u>AB2</u>	MYLAN PHARMS INC	<u>500MG</u>	<u>A200690 001</u>	Aug 01, 2012
<u>AB2</u>		<u>1GM</u>	<u>A200690 002</u>	Aug 01, 2012
<u>AB2</u>	NOSTRUM LABS INC	<u>500MG</u>	<u>A203832 001</u>	Dec 26, 2017
<u>AB2</u>		<u>1GM</u>	<u>A203832 002</u>	Dec 26, 2017
<u>AB2</u>	NOVAST LABS	<u>500MG</u>	<u>A209674 001</u>	Nov 02, 2018
<u>AB2</u>		<u>1GM</u>	<u>A209674 002</u>	Nov 02, 2018
<u>AB2</u>	QINGDAO BAHEAL PHARM	<u>500MG</u>	<u>A209993 001</u>	Dec 27, 2018
<u>AB2</u>		<u>1GM</u>	<u>A209993 002</u>	Dec 27, 2018

GLUMETZA

<u>AB3</u>	+ SANTARUS INC	<u>500MG</u>	<u>N021748 001</u>	Jun 03, 2005
<u>AB3</u>	+	<u>1GM</u>	<u>N021748 002</u>	Jun 03, 2005

METFORMIN HYDROCHLORIDE

<u>AB3</u>	ACTAVIS LABS FL INC	<u>500MG</u>	<u>A203755 001</u>	Aug 01, 2016
<u>AB3</u>		<u>1GM</u>	<u>A203755 002</u>	Aug 01, 2016
<u>AB3</u>	GLENMARK PHARMS LTD	<u>500MG</u>	<u>A212969 001</u>	Nov 25, 2019
<u>AB3</u>		<u>1GM</u>	<u>A212969 002</u>	Nov 25, 2019
<u>AB3</u>	LUPIN LTD	<u>500MG</u>	<u>A091664 001</u>	Jul 19, 2013
<u>AB3</u>		<u>1GM</u>	<u>A091664 002</u>	Jul 19, 2013
<u>AB3</u>	SUN PHARM	<u>500MG</u>	<u>A202917 001</u>	Aug 01, 2016
<u>AB3</u>		<u>1GM</u>	<u>A202917 002</u>	Aug 01, 2016

METFORMIN HYDROCHLORIDE; PIOGLITAZONE HYDROCHLORIDE

TABLET;ORAL

ACTOPLUS MET

<u>AB</u>	+ TAKEDA PHARMS USA	<u>500MG;EQ 15MG BASE</u>	<u>N021842 001</u>	Aug 29, 2005
<u>AB</u>	+	<u>850MG;EQ 15MG BASE</u>	<u>N021842 002</u>	Aug 29, 2005

PIOGLITAZONE HYDROCHLORIDE AND METFORMIN HYDROCHLORIDE

<u>AB</u>	AUROBINDO PHARMA LTD	<u>500MG;EQ 15MG BASE</u>	<u>A200823 001</u>	Feb 13, 2013
<u>AB</u>		<u>850MG;EQ 15MG BASE</u>	<u>A200823 002</u>	Feb 13, 2013
<u>AB</u>	MACLEODS PHARMS LTD	<u>500MG;EQ 15MG BASE</u>	<u>A204802 001</u>	Nov 05, 2015
<u>AB</u>		<u>850MG;EQ 15MG BASE</u>	<u>A204802 002</u>	Nov 05, 2015
<u>AB</u>	SANDOZ	<u>500MG;EQ 15MG BASE</u>	<u>A091273 001</u>	Apr 16, 2013
<u>AB</u>		<u>850MG;EQ 15MG BASE</u>	<u>A091273 002</u>	Apr 16, 2013
<u>AB</u>	TEVA PHARMS USA	<u>500MG;EQ 15MG BASE</u>	<u>A091155 001</u>	Mar 10, 2014
<u>AB</u>		<u>850MG;EQ 15MG BASE</u>	<u>A091155 002</u>	Mar 10, 2014
<u>AB</u>	TORRENT PHARMS LTD	<u>500MG;EQ 15MG BASE</u>	<u>A202001 001</u>	Feb 13, 2013
<u>AB</u>		<u>850MG;EQ 15MG BASE</u>	<u>A202001 002</u>	Feb 13, 2013

METFORMIN HYDROCHLORIDE; SAXAGLIPTIN HYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL

KOMBIGLYZE XR

+	ASTRAZENECA AB	<u>500MG;EQ 5MG BASE</u>	N200678 001	Nov 05, 2010
+		<u>1GM;EQ 2.5MG BASE</u>	N200678 003	Nov 05, 2010
+	!	<u>1GM;EQ 5MG BASE</u>	N200678 002	Nov 05, 2010

METFORMIN HYDROCHLORIDE; SITAGLIPTIN PHOSPHATE

TABLET;ORAL

JANUMET

+	MERCK SHARP DOHME	<u>500MG;EQ 50MG BASE</u>	N022044 001	Mar 30, 2007
+	!	<u>1GM;EQ 50MG BASE</u>	N022044 002	Mar 30, 2007

TABLET, EXTENDED RELEASE;ORAL

JANUMET XR

+	MERCK SHARP DOHME	<u>500MG;EQ 50MG BASE</u>	N202270 001	Feb 02, 2012
+		<u>1GM;EQ 50MG BASE</u>	N202270 002	Feb 02, 2012
+	!	<u>1GM;EQ 100MG BASE</u>	N202270 003	Feb 02, 2012

METHACHOLINE CHLORIDE

FOR SOLUTION;INHALATION

PROVOCHOLINE

+	METHAPHARM	<u>100MG/VIAL</u>	N019193 001	Oct 31, 1986
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PRESCRIPTION DRUG PRODUCT LIST

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>	PHARM ASSOC	<u>10MG/ML</u>	<u>A207368</u>	<u>001</u>	Aug 22, 2019
<u>AA</u>	VISTAPHARM	<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
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METHADOSE

<u>AA</u>	+!	SPECGX LLC	<u>10MG/ML</u>	<u>N017116</u>	<u>002</u>
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INJECTABLE; INJECTION

METHADONE HYDROCHLORIDE

<u>AP</u>		AKORN	<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>		PHARM ASSOC	<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
<u>AA</u>		VISTAPHARM	<u>5MG/5ML</u>	<u>A090707</u>	<u>001</u>	Jun 30, 2010
<u>AA</u>			<u>10MG/5ML</u>	<u>A090707</u>	<u>002</u>	Jun 30, 2010

TABLET; ORAL

DOLOPHINE HYDROCHLORIDE

<u>AA</u>	+!	HIKMA	<u>5MG</u>	<u>N006134</u>	<u>002</u>
<u>AA</u>	+!		<u>10MG</u>	<u>N006134</u>	<u>010</u>

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>		SPECGX LLC	<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>		SUN PHARM	<u>5MG</u>	<u>A208305</u>	<u>001</u>	Mar 30, 2018
<u>AA</u>		INDUSTRIES				
<u>AA</u>			<u>10MG</u>	<u>A208305</u>	<u>002</u>	Mar 30, 2018
<u>AA</u>		THEPHARMANETWORK	<u>10MG</u>	<u>A090635</u>	<u>001</u>	Nov 25, 2009
<u>AA</u>		LLC				
<u>AA</u>		VISTAPHARM	<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

METHADOSE

<u>AA</u>		SPECGX LLC	<u>5MG</u>	<u>A040050</u>	<u>001</u>	Apr 15, 1993
<u>AA</u>			<u>10MG</u>	<u>A040050</u>	<u>002</u>	Apr 15, 1993

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

METHADOSE

<u>AA</u>		SPECGX LLC	<u>40MG</u>	<u>A074184</u>	<u>001</u>	Apr 29, 1993
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METHAMPHETAMINE HYDROCHLORIDE

TABLET; ORAL

DESOXYN

<u>AA</u>	+!	RECORDATI RARE	<u>5MG</u>	<u>N005378</u>	<u>002</u>
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METHAMPHETAMINE HYDROCHLORIDE

<u>AA</u>		HIKMA	<u>5MG</u>	<u>A203846</u>	<u>001</u>	Nov 17, 2015
<u>AA</u>		MAYNE PHARMA INC	<u>5MG</u>	<u>A091189</u>	<u>001</u>	Apr 21, 2010

METHAZOLAMIDE

TABLET; ORAL

METHAZOLAMIDE

<u>AB</u>		ANI PHARMS INC	<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>		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>		SANDOZ	<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

PRESCRIPTION DRUG PRODUCT LIST

METHENAMINE HIPPURATE

TABLET; ORAL

HIPREX

<u>AB</u>	<u>+!</u> US PHARM HOLDINGS	<u>1GM</u>	<u>N017681</u>	<u>001</u>	
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METHENAMINE HIPPURATE

<u>AB</u>	AUROBINDO PHARMA LTD	<u>1GM</u>	<u>A205661</u>	<u>001</u>	Jul 05, 2016
<u>AB</u>	IMPAX LABS INC	<u>1GM</u>	<u>A076411</u>	<u>001</u>	Jun 20, 2003
<u>AB</u>	MICRO LABS	<u>1GM</u>	<u>A212172</u>	<u>001</u>	Aug 01, 2019

UREX

<u>AB</u>	CNTY LINE PHARMS	<u>1GM</u>	<u>N016151</u>	<u>001</u>	
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METHIMAZOLE

TABLET; ORAL

METHIMAZOLE

<u>AB</u>	CASI PHARMS INC	<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>	ECI PHARMS LLC	<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>	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>	MYLAN	<u>5MG</u>	<u>A040350</u>	<u>001</u>	Mar 29, 2000
<u>AB</u>	<u>!</u>	<u>10MG</u>	<u>A040350</u>	<u>002</u>	Mar 29, 2000
<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

TAPAZOLE

<u>AB</u>	KING PHARMS LLC	<u>5MG</u>	<u>A040320</u>	<u>001</u>	Mar 31, 2000
<u>AB</u>		<u>10MG</u>	<u>A040320</u>	<u>002</u>	Mar 31, 2000

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>	AUROBINDO PHARMA LTD	<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>	MYLAN INSTITUTIONAL	<u>1GM/10ML (100MG/ML)</u>	<u>A204404</u>	<u>001</u>	Dec 05, 2014
<u>AP</u>	NAVINTA LLC	<u>1GM/10ML (100MG/ML)</u>	<u>A206071</u>	<u>001</u>	Nov 24, 2017
<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	<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>	<u>+!</u> HIKMA	<u>1GM/10ML (100MG/ML)</u>	<u>N011790</u>	<u>001</u>	
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TABLET; ORAL

METHOCARBAMOL

<u>AA</u>	AUSTARPHARMA LLC	<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>	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>	DBL PHARMS	<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>	<u>!</u> GRANULES INDIA LTD	<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>	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>	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>A086989</u>	<u>001</u>	
<u>AA</u>		<u>750MG</u>	<u>A086988</u>	<u>001</u>	

ROBAXIN-750

<u>AA</u>	<u>+!</u> AUXILIUM PHARMS LLC	<u>750MG</u>	<u>N011011</u>	<u>006</u>	
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METHOHEXITAL SODIUM

INJECTABLE; INJECTION

BREVITAL SODIUM

<u>+!</u>	PAR STERILE PRODUCTS	500MG/VIAL	N011559	001	
<u>+!</u>		2.5GM/VIAL	N011559	002	

PRESCRIPTION DRUG PRODUCT LIST

3-289 (of 453)

METHOTREXATE

SOLUTION; SUBCUTANEOUS

OTREXUP

+	!	ANTARES PHARMA INC	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

REDITREX

CUMBERLAND PHARMS

			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

METHOTREXATE PRESERVATIVE FREE

<u>AP</u>		FRESENIUS KABI USA	<u>EQ 25MG BASE/ML</u>	<u>A040265</u>	<u>001</u>	Feb 26, 1999
<u>AP</u>			<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>	+	HOSPIRA	<u>EQ 50MG BASE/2ML (EQ 25MG BASE/ML)</u>	<u>N011719</u>	<u>010</u>	Dec 15, 2004
<u>AP</u>	!	WEST-WARD PHARMS INT	<u>EQ 100MG BASE/4ML (EQ 25MG BASE/ML)</u>	<u>A089341</u>	<u>001</u>	Sep 16, 1986

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>	+	HOSPIRA	<u>EQ 1GM BASE/40ML (EQ 25MG BASE/ML)</u>	<u>N011719</u>	<u>012</u>	Apr 13, 2005
<u>AP</u>		MYLAN LABS LTD	<u>EQ 250MG BASE/10ML (EQ 25MG BASE/ML)</u>	<u>A201529</u>	<u>004</u>	Mar 29, 2012
<u>AP</u>			<u>EQ 1GM BASE/40ML (EQ 25MG BASE/ML)</u>	<u>A201530</u>	<u>001</u>	Mar 29, 2012
<u>AP</u>		PHARMACHEMIE BV	<u>EQ 50MG BASE/2ML (EQ 25MG BASE/ML)</u>	<u>A040843</u>	<u>002</u>	Jan 11, 2010
<u>AP</u>			<u>EQ 250MG BASE/10ML (EQ 25MG BASE/ML)</u>	<u>A040843</u>	<u>004</u>	Jan 11, 2010
<u>AP</u>			<u>EQ 1GM BASE/40ML (EQ 25MG BASE/ML)</u>	<u>A040843</u>	<u>001</u>	Jan 11, 2010
<u>AP</u>		SAGENT PHARMS INC	<u>EQ 50MG BASE/2ML (EQ 25MG BASE/ML)</u>	<u>A203407</u>	<u>001</u>	Aug 09, 2018
<u>AP</u>			<u>EQ 250MG BASE/10ML (EQ 25MG BASE/ML)</u>	<u>A203407</u>	<u>002</u>	Aug 09, 2018
<u>AP</u>			<u>EQ 1GM BASE/40ML (EQ 25MG BASE/ML)</u>	<u>A203407</u>	<u>003</u>	Aug 09, 2018
<u>AP</u>		SANDOZ INC	<u>EQ 50MG BASE/2ML (EQ 25MG BASE/ML)</u>	<u>A090039</u>	<u>001</u>	Mar 31, 2009
<u>AP</u>			<u>EQ 250MG BASE/10ML (EQ 25MG BASE/ML)</u>	<u>A090039</u>	<u>002</u>	Mar 31, 2009
<u>AP</u>			<u>EQ 1GM BASE/40ML (EQ 25MG BASE/ML)</u>	<u>A090029</u>	<u>001</u>	Mar 31, 2009
<u>AP</u>	!	WEST-WARD PHARMS INT	<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

METHOTREXATE SODIUM

!		WEST-WARD PHARMS INT	EQ 200MG BASE/8ML (EQ 25MG BASE/ML)	A089342	001	Sep 16, 1986
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METHOTREXATE SODIUM PRESERVATIVE FREE

		PHARMACHEMIE BV	EQ 100MG BASE/4ML (EQ 25MG BASE/ML)	A040843	003	Feb 27, 2012
!		WEST-WARD PHARMS INT	EQ 1GM BASE/VIAL	A040632	001	Aug 12, 2005

SOLUTION; ORAL

XATMEP

+	!	SILVERGATE PHARMS	EQ 2MG BASE/ML	N208400	001	Apr 25, 2017
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TABLET; ORAL

METHOTREXATE SODIUM

<u>AB</u>		ACCORD HLTHCARE	<u>EQ 2.5MG BASE</u>	<u>A213343</u>	<u>001</u>	Jan 24, 2020
<u>AB</u>		AMNEAL PHARMS	<u>EQ 2.5MG BASE</u>	<u>A210040</u>	<u>001</u>	Dec 22, 2017
<u>AB</u>		BARR	<u>EQ 2.5MG BASE</u>	<u>A081099</u>	<u>001</u>	Oct 15, 1990

PRESCRIPTION DRUG PRODUCT LIST

METHOTREXATE SODIUM

TABLET; ORAL

METHOTREXATE SODIUM

<u>AB</u>	+ !	DAVA PHARMS INC	<u>EQ 2.5MG BASE</u>	<u>N008085 002</u>	
<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

METHOXSALEN

CAPSULE; ORAL

METHOXSALEN

<u>AB</u>		ACTAVIS INC	<u>10MG</u>	<u>A202603 001</u>	Jun 09, 2015
<u>AB</u>		STRIDES PHARMA	<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			
		+ !	MALLINCKRODT HOSP	0.02MG/ML	N020969 001 Feb 25, 1999

METHSCOPOLAMINE BROMIDE

TABLET; ORAL

METHSCOPOLAMINE BROMIDE

		BAYSHORE PHARMS LLC	2.5MG	A200602 001	Sep 24, 2012
		!	5MG	A200602 002	Sep 24, 2012

METHSUXIMIDE

CAPSULE; ORAL

CELONTIN

		+ !	PARKE DAVIS	300MG	N010596 008
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METHYLDOPA

TABLET; ORAL

METHYLDOPA

<u>AB</u>		ACCORD HLTHCARE	<u>250MG</u>	<u>A070084 001</u>	Oct 15, 1985
<u>AB</u>			<u>500MG</u>	<u>A070085 001</u>	Oct 15, 1985
<u>AB</u>		HERITAGE PHARMA	<u>250MG</u>	<u>A070098 001</u>	Feb 20, 1986
<u>AB</u>			<u>500MG</u>	<u>A070343 001</u>	Feb 20, 1986
<u>AB</u>		MYLAN	<u>250MG</u>	<u>A070076 002</u>	Apr 18, 1985
<u>AB</u>	!		<u>500MG</u>	<u>A070076 001</u>	Apr 18, 1985
<u>AB</u>		WATSON LABS	<u>500MG</u>	<u>A070625 001</u>	Jun 06, 1986

METHYLENE BLUE

SOLUTION; INTRAVENOUS

PROVAYBLUE

		+ !	PROVEPHARM SAS	50MG/10ML (5MG/ML)	N204630 001 Apr 08, 2016
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METHYLERGONOVINE MALEATE

INJECTABLE; INJECTION

METHERGINE

<u>AP</u>	+ !	EDISON THERAPS LLC	<u>0.2MG/ML</u>	<u>N006035 004</u>	
		<u>METHYLERGONOVINE MALEATE</u>			
<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>		GRANULES PHARMS	<u>0.2MG</u>	<u>A210424 001</u>	May 15, 2018
<u>AB</u>	!	NOVEL LABS INC	<u>0.2MG</u>	<u>A091577 001</u>	May 02, 2011
<u>AB</u>		TEVA PHARMS USA	<u>0.2MG</u>	<u>A211455 001</u>	Mar 20, 2019

METHYLNALTREXONE BROMIDE

SOLUTION; SUBCUTANEOUS

RELISTOR

		+ !	SALIX PHARMS	8MG/0.4ML (8MG/0.4ML)	N021964 002 Sep 27, 2010
		+ !		12MG/0.6ML (12MG/0.6ML)	N021964 001 Apr 24, 2008
		+ !		12MG/0.6ML (12MG/0.6ML)	N021964 003 Sep 27, 2010

PRESCRIPTION DRUG PRODUCT LIST

METHYLNALTREXONE BROMIDE

TABLET;ORAL

RELISTOR

+! SALIX 150MG N208271 001 Jul 19, 2016

METHYLPHENIDATE

FILM, EXTENDED RELEASE;TRANSDERMAL

DAYTRANA

+ NOVEN PHARMS INC 10MG/9HR (1.1MG/HR) N021514 001 Apr 06, 2006
 + 15MG/9HR (1.6MG/HR) N021514 002 Apr 06, 2006
 + 20MG/9HR (2.2MG/HR) N021514 003 Apr 06, 2006
 +! 30MG/9HR (3.3MG/HR) N021514 004 Apr 06, 2006

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

AB1 BARR LABS INC 10MG A079031 004 Oct 15, 2014
AB1 20MG A079031 001 Jul 13, 2012
AB1 30MG A079031 002 Jul 13, 2012
AB1 40MG A079031 003 Jul 13, 2012
AB1 GRANULES PHARMS 10MG A211796 001 May 23, 2019
AB1 20MG A211796 002 May 23, 2019
AB1 30MG A211796 003 May 23, 2019
AB1 40MG A211796 004 May 23, 2019
AB1 60MG A211796 005 May 23, 2019
AB1 MAYNE PHARMA 10MG A200886 001 Feb 26, 2018
AB1 20MG A078458 001 Dec 01, 2011
AB1 30MG A078458 002 Dec 01, 2011
AB1 40MG A078458 003 Dec 01, 2011
AB1 ! 60MG A078458 004 Jun 23, 2016

RITALIN LA

AB1 + NOVARTIS 10MG N021284 004 Apr 10, 2004
AB1 + 20MG N021284 001 Jun 05, 2002
AB1 + 30MG N021284 002 Jun 05, 2002
AB1 + 40MG N021284 003 Jun 05, 2002

METADATE CD

AB2 + LANNETT CO INC 10MG N021259 003 May 27, 2003
AB2 + 20MG N021259 001 Apr 03, 2001
AB2 + 30MG N021259 002 Jun 19, 2003
AB2 + 40MG N021259 004 Feb 19, 2006
AB2 + 50MG N021259 005 Feb 19, 2006
AB2 +! 60MG N021259 006 Feb 19, 2006

METHYLPHENIDATE HYDROCHLORIDE

AB2 IMPAX LABS INC 10MG A205105 001 Jul 28, 2016
AB2 20MG A205105 002 Jul 28, 2016
AB2 30MG A205105 003 Jul 28, 2016
AB2 40MG A205105 004 Jul 28, 2016
AB2 50MG A205105 005 Jul 28, 2016
AB2 60MG A205105 006 Jul 28, 2016
AB2 SPECGX LLC 10MG A203583 001 Sep 29, 2015
AB2 20MG A203583 002 Sep 29, 2015
AB2 30MG A203583 003 Sep 29, 2015
AB2 40MG A203583 004 Sep 29, 2015
AB2 50MG A203583 005 Sep 29, 2015
AB2 60MG A203583 006 Sep 29, 2015
AB2 TEVA PHARMS 10MG A077707 001 Jul 19, 2012
AB2 20MG A077707 002 Jul 19, 2012
AB2 30MG A077707 003 Jul 19, 2012
AB2 40MG A078873 001 Jul 19, 2012
AB2 50MG A078873 002 Jul 19, 2012
AB2 60MG A078873 003 Jul 19, 2012

APTENSIO XR

AB3 + RHODES PHARMS 10MG N205831 001 Apr 17, 2015
AB3 + 15MG N205831 002 Apr 17, 2015
AB3 + 20MG N205831 003 Apr 17, 2015
AB3 + 30MG N205831 004 Apr 17, 2015
AB3 + 40MG N205831 005 Apr 17, 2015
AB3 + 50MG N205831 006 Apr 17, 2015

PRESCRIPTION DRUG PRODUCT LIST

METHYLPHENIDATE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE;ORAL

APTENSIO XRAB3 +! 60MG N205831 007 Apr 17, 2015METHYLPHENIDATE HYDROCHLORIDEAB3 ACTAVIS ELIZABETH 10MG A208861 001 Dec 13, 2018AB3 15MG A208861 002 Dec 13, 2018AB3 20MG A208861 003 Dec 13, 2018AB3 30MG A208861 004 Dec 13, 2018AB3 40MG A208861 005 Dec 13, 2018AB3 50MG A208861 006 Dec 13, 2018AB3 60MG A208861 007 Dec 13, 2018

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

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

QUILLIVANT XR

+! NEXTWAVE 5MG/ML N202100 001 Sep 27, 2012

SOLUTION;ORAL

METHYLINAA +! SPECGX LLC 5MG/5ML N021419 001 Dec 19, 2002AA +! 10MG/5ML N021419 002 Dec 19, 2002METHYLPHENIDATE HYDROCHLORIDEAA ABHAI LLC 5MG/5ML A207485 001 Nov 18, 2016AA 10MG/5ML A207485 002 Nov 18, 2016AA BRECKENRIDGE 5MG/5ML A201466 001 Nov 12, 2013AA 10MG/5ML A201466 002 Nov 12, 2013AA NOVEL LABS INC 5MG/5ML A204602 001 Aug 14, 2015AA 10MG/5ML A204602 002 Aug 14, 2015AA NOVELGENIX THERAPS 5MG/5ML A210139 001 Oct 03, 2018AA 10MG/5ML A210139 002 Oct 03, 2018AA TRIS PHARMA INC 5MG/5ML A091601 001 Jul 23, 2010AA 10MG/5ML A091601 002 Jul 23, 2010

TABLET;ORAL

METHYLPHENIDATE HYDROCHLORIDEAB ABHAI INC 5MG A206932 001 May 11, 2017AB 10MG A206932 002 May 11, 2017AB 20MG A206932 003 May 11, 2017AB ACTAVIS LABS FL INC 5MG A040220 001 Aug 29, 1997AB 10MG A040220 002 Aug 29, 1997AB 20MG A040220 003 Aug 29, 1997AB ALKEM LABS LTD 5MG A211779 001 Oct 04, 2019AB 10MG A211779 002 Oct 04, 2019AB 20MG A211779 003 Oct 04, 2019AB ASCENT PHARMS INC 5MG A207416 001 Sep 22, 2015AB 10MG A207416 002 Sep 22, 2015AB 20MG A207416 003 Sep 22, 2015AB BIONPHARMA INC 5MG A209753 001 Mar 02, 2018AB 10MG A209753 002 Mar 02, 2018AB 20MG A209753 003 Mar 02, 2018AB BRECKENRIDGE 5MG A207587 001 Mar 03, 2017AB 10MG A207587 002 Mar 03, 2017AB 20MG A207587 003 Mar 03, 2017AB MOUNTAIN 5MG A091159 001 Mar 12, 2014AB 10MG A091159 002 Mar 12, 2014AB 20MG A091159 003 Mar 12, 2014AB NOVEL LABS INC 5MG A207884 001 Nov 13, 2015AB 10MG A207884 002 Nov 13, 2015AB 20MG A207884 003 Nov 13, 2015AB OXFORD PHARMS 5MG A202892 001 Sep 23, 2014AB 10MG A202892 002 Sep 23, 2014AB 20MG A202892 003 Sep 23, 2014

PRESCRIPTION DRUG PRODUCT LIST

METHYLPHENIDATE HYDROCHLORIDE

TABLET; ORAL

METHYLPHENIDATE HYDROCHLORIDE

<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>	+	NOVARTIS	<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>	BRECKENRIDGE	<u>2.5MG</u>	<u>A204954 001</u>	Jan 26, 2017
<u>AB</u>		<u>5MG</u>	<u>A204954 002</u>	Jan 26, 2017
<u>AB</u>		<u>10MG</u>	<u>A204954 003</u>	Jan 26, 2017
<u>AB</u>	NOVEL LABS INC	<u>2.5MG</u>	<u>A204115 001</u>	Feb 25, 2015
<u>AB</u>		<u>5MG</u>	<u>A204115 002</u>	Feb 25, 2015
<u>AB</u>	!	<u>10MG</u>	<u>A204115 003</u>	Feb 25, 2015
<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
<u>AB</u>	ALVOGEN PINE BROOK	<u>18MG</u>	<u>A210818 001</u>	Nov 30, 2018
<u>AB</u>		<u>27MG</u>	<u>A210818 002</u>	Nov 30, 2018
<u>AB</u>		<u>36MG</u>	<u>A210818 003</u>	Nov 30, 2018
<u>AB</u>		<u>54MG</u>	<u>A210818 004</u>	Nov 30, 2018
<u>AB</u>	AMNEAL PHARMS	<u>18MG</u>	<u>A207515 001</u>	Feb 01, 2018
<u>AB</u>		<u>27MG</u>	<u>A207515 002</u>	Feb 01, 2018
<u>AB</u>		<u>36MG</u>	<u>A207515 003</u>	Feb 01, 2018
<u>AB</u>		<u>54MG</u>	<u>A207515 004</u>	Feb 01, 2018
<u>AB</u>	ANDOR PHARMS	<u>18MG</u>	<u>A211918 001</u>	Apr 24, 2019
<u>AB</u>		<u>27MG</u>	<u>A211918 002</u>	Apr 24, 2019
<u>AB</u>		<u>36MG</u>	<u>A211918 003</u>	Apr 24, 2019
<u>AB</u>		<u>54MG</u>	<u>A211918 004</u>	Apr 24, 2019
<u>AB</u>	ANI PHARMS INC	<u>18MG</u>	<u>A208607 001</u>	Jul 14, 2017
<u>AB</u>		<u>27MG</u>	<u>A208607 002</u>	Jul 14, 2017
<u>AB</u>		<u>36MG</u>	<u>A208607 003</u>	Jul 14, 2017
<u>AB</u>		<u>54MG</u>	<u>A208607 004</u>	Jul 14, 2017
<u>AB</u>	ASCENT PHARMS INC	<u>18MG</u>	<u>A211009 001</u>	Sep 03, 2019
<u>AB</u>		<u>27MG</u>	<u>A211009 002</u>	Sep 03, 2019
<u>AB</u>		<u>36MG</u>	<u>A211009 003</u>	Sep 03, 2019
<u>AB</u>		<u>54MG</u>	<u>A211009 004</u>	Sep 03, 2019
<u>AB</u>	CNTY LINE PHARMS	<u>10MG</u>	<u>A204772 001</u>	Feb 29, 2016
<u>AB</u>		<u>20MG</u>	<u>A204772 002</u>	Feb 29, 2016
<u>AB</u>	GRANULES PHARMS	<u>10MG</u>	<u>A210992 001</u>	Nov 21, 2018
<u>AB</u>		<u>20MG</u>	<u>A210992 002</u>	Nov 21, 2018
<u>AB</u>	MYLAN	<u>18MG</u>	<u>A206726 001</u>	Oct 21, 2016
<u>AB</u>		<u>27MG</u>	<u>A206726 002</u>	Oct 21, 2016
<u>AB</u>		<u>36MG</u>	<u>A206726 003</u>	Oct 21, 2016
<u>AB</u>		<u>54MG</u>	<u>A206726 004</u>	Oct 21, 2016
<u>AB</u>	OSMOTICA	<u>18MG</u>	<u>A205327 001</u>	Jul 28, 2017

PRESCRIPTION DRUG PRODUCT LIST

METHYLPHENIDATE HYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL

METHYLPHENIDATE HYDROCHLORIDE

<u>AB</u>		<u>27MG</u>	<u>A205327 002</u>	Jul 28, 2017
<u>AB</u>		<u>36MG</u>	<u>A205327 003</u>	Jul 28, 2017
<u>AB</u>		<u>54MG</u>	<u>A205327 004</u>	Jul 28, 2017
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
!	OSMOTICA	72MG	A205327 005	Jul 28, 2017

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

<u>AB</u>	+	PHARMACIA AND UPJOHN	<u>2MG</u>	<u>N011153 002</u>
<u>AB</u>	+		<u>4MG</u>	<u>N011153 001</u>
<u>AB</u>	+		<u>8MG</u>	<u>N011153 004</u>
<u>AB</u>	+		<u>16MG</u>	<u>N011153 003</u>
<u>AB</u>	+	!	<u>32MG</u>	<u>N011153 006</u>

METHYLPREDNISOLONE

<u>AB</u>	DURAMED PHARMS BARR	<u>4MG</u>	<u>A088497 001</u>	Feb 21, 1984
<u>AB</u>	JUBILANT CADISTA	<u>4MG</u>	<u>A040189 001</u>	Oct 31, 1997
<u>AB</u>		<u>8MG</u>	<u>A040189 002</u>	Oct 31, 1997
<u>AB</u>		<u>16MG</u>	<u>A040189 003</u>	Jul 20, 2007
<u>AB</u>		<u>32MG</u>	<u>A040189 004</u>	Jul 20, 2007
<u>AB</u>	SANDOZ	<u>4MG</u>	<u>A040194 001</u>	Oct 31, 1997
<u>AB</u>	SUNGEN PHARMA	<u>4MG</u>	<u>A212262 001</u>	Jun 27, 2019
<u>AB</u>	TIANJIN TIANYAO	<u>4MG</u>	<u>A204072 001</u>	May 14, 2018
<u>AB</u>	VINTAGE PHARMS	<u>4MG</u>	<u>A040183 001</u>	Dec 22, 1998
<u>AB</u>	WATSON LABS	<u>4MG</u>	<u>A040232 001</u>	Oct 16, 1997
<u>AB</u>	ZYDUS PHARMS	<u>4MG</u>	<u>A206751 001</u>	Apr 23, 2018
<u>AB</u>		<u>8MG</u>	<u>A206751 002</u>	Apr 23, 2018
<u>AB</u>		<u>16MG</u>	<u>A206751 003</u>	Apr 23, 2018
<u>AB</u>		<u>32MG</u>	<u>A206751 004</u>	Apr 23, 2018

METHYLPREDNISOLONE ACETATE

INJECTABLE; INJECTION

DEPO-MEDROL

<u>AB</u>	+	PHARMACIA AND UPJOHN	<u>20MG/ML</u>	<u>N011757 002</u>
<u>AB</u>	+		<u>40MG/ML</u>	<u>N011757 001</u>
<u>AB</u>	+		<u>80MG/ML</u>	<u>N011757 004</u>

METHYLPREDNISOLONE ACETATE

<u>AB</u>	SAGENT PHARMS INC	<u>20MG/ML</u>	<u>A201835 001</u>	Jun 27, 2018
<u>AB</u>		<u>40MG/ML</u>	<u>A201835 002</u>	Jun 27, 2018
<u>AB</u>		<u>80MG/ML</u>	<u>A201835 003</u>	Jun 27, 2018
<u>AB</u>	SANDOZ INC	<u>40MG/ML</u>	<u>A040719 001</u>	Jan 29, 2009
<u>AB</u>		<u>40MG/ML</u>	<u>A040794 001</u>	Mar 05, 2009
<u>AB</u>		<u>80MG/ML</u>	<u>A040719 002</u>	Jan 29, 2009
<u>AB</u>		<u>80MG/ML</u>	<u>A040794 002</u>	Mar 05, 2009
<u>AB</u>	TEVA PHARMS USA	<u>40MG/ML</u>	<u>A040557 001</u>	Feb 23, 2005
<u>AB</u>		<u>40MG/ML</u>	<u>A040620 001</u>	Oct 27, 2006
<u>AB</u>		<u>80MG/ML</u>	<u>A040557 002</u>	Feb 23, 2005
<u>AB</u>		<u>80MG/ML</u>	<u>A040620 002</u>	Oct 27, 2006

METHYLPREDNISOLONE SODIUM SUCCINATE

INJECTABLE; INJECTION

METHYLPREDNISOLONE SODIUM SUCCINATE

<u>AP</u>	AMNEAL	<u>EQ 40MG BASE/VIAL</u>	<u>A207549 001</u>	Nov 09, 2016
<u>AP</u>		<u>EQ 125MG BASE/VIAL</u>	<u>A207549 002</u>	Nov 09, 2016
<u>AP</u>	AUROBINDO PHARMA LTD	<u>EQ 40MG BASE/VIAL</u>	<u>A207667 001</u>	Dec 15, 2015
<u>AP</u>		<u>EQ 125MG BASE/VIAL</u>	<u>A207667 002</u>	Dec 15, 2015
<u>AP</u>		<u>EQ 500MG BASE/VIAL</u>	<u>A207667 003</u>	Dec 15, 2015

PRESCRIPTION DRUG PRODUCT LIST

METHYLPREDNISOLONE SODIUM SUCCINATE

INJECTABLE; INJECTION

METHYLPREDNISOLONE SODIUM SUCCINATE

<u>AP</u>		<u>EQ 2GM BASE/VIAL</u>	<u>A207667 004</u>	Dec 15, 2015
<u>AP</u>	FRESENIUS KABI USA	<u>EQ 40MG BASE/VIAL</u>	<u>A040583 001</u>	Jul 30, 2004
<u>AP</u>		<u>EQ 125MG BASE/VIAL</u>	<u>A040583 002</u>	Jul 30, 2004
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>A040612 001</u>	Aug 12, 2004
<u>AP</u>	HIKMA	<u>EQ 500MG BASE/VIAL</u>	<u>A202691 001</u>	Feb 16, 2016
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>A202691 002</u>	Feb 16, 2016
<u>AP</u>	SAGENT PHARMS INC	<u>EQ 40MG BASE/VIAL</u>	<u>A040888 001</u>	Jul 18, 2011
<u>AP</u>		<u>EQ 125MG BASE/VIAL</u>	<u>A040888 002</u>	Jul 18, 2011
<u>AP</u>		<u>EQ 500MG BASE/VIAL</u>	<u>A040888 003</u>	Jul 18, 2011
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>A040888 004</u>	Jul 18, 2011
<u>AP</u>		<u>EQ 2GM BASE/VIAL</u>	<u>A040888 005</u>	Jul 18, 2011
<u>SOLU-MEDROL</u>				
<u>AP</u>	+! PHARMACIA AND UPJOHN	<u>EQ 40MG BASE/VIAL</u>	<u>N011856 003</u>	
<u>AP</u>	+!	<u>EQ 125MG BASE/VIAL</u>	<u>N011856 004</u>	
<u>AP</u>	+!	<u>EQ 500MG BASE/VIAL</u>	<u>N011856 005</u>	
<u>AP</u>	+!	<u>EQ 1GM BASE/VIAL</u>	<u>N011856 006</u>	
<u>AP</u>	+!	<u>EQ 2GM BASE/VIAL</u>	<u>N011856 007</u>	Feb 27, 1985

METHYLTESTOSTERONE

CAPSULE; ORAL

METHYLTESTOSTERONE

<u>AB</u>	IMPAX LABS INC	<u>10MG</u>	<u>A204851 001</u>	Sep 21, 2015
<u>TESTRED</u>				
<u>AB</u>	! VALEANT PHARM INTL	<u>10MG</u>	<u>A083976 001</u>	
TABLET; ORAL				
ANDROID 25				
<u>BP</u>	VALEANT PHARM INTL	25MG	A087147 001	
METHYLTESTOSTERONE				
<u>BP</u>	IMPAX LABS	10MG	A080767 002	

METOCLOPRAMIDE HYDROCHLORIDE

INJECTABLE; INJECTION

METOCLOPRAMIDE

<u>AP</u>	EMCURE PHARMS LTD	<u>EQ 5MG BASE/ML</u>	<u>A204756 001</u>	Dec 20, 2013
<u>METOCLOPRAMIDE HYDROCHLORIDE</u>				
<u>AP</u>	FRESENIUS KABI USA	<u>EQ 5MG BASE/ML</u>	<u>A091392 001</u>	Apr 19, 2013
<u>AP</u>	! HOSPIRA	<u>EQ 5MG BASE/ML</u>	<u>A073118 001</u>	Jan 17, 1991
<u>AP</u>	TEVA PHARMS USA	<u>EQ 5MG BASE/ML</u>	<u>A073135 001</u>	Nov 27, 1991

SOLUTION; ORAL

METOCLOPRAMIDE HYDROCHLORIDE

<u>AA</u>	ANI PHARMS	<u>EQ 5MG BASE/5ML</u>	<u>A071402 001</u>	Jun 25, 1993
<u>AA</u>	PHARM ASSOC	<u>EQ 5MG BASE/5ML</u>	<u>A072744 001</u>	May 28, 1991
<u>AA</u>	VISTAPHARM	<u>EQ 5MG BASE/5ML</u>	<u>A075051 001</u>	Jan 26, 2001
<u>AA</u>	! WOCKHARDT BIO AG	<u>EQ 5MG BASE/5ML</u>	<u>A074703 001</u>	Oct 31, 1997

TABLET; ORAL

METOCLOPRAMIDE HYDROCHLORIDE

<u>AB</u>	IMPAX LABS INC	<u>EQ 5MG BASE</u>	<u>A071250 002</u>	Dec 28, 1995
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A071250 001</u>	Feb 03, 1988
<u>AB</u>	IPCA LABS LTD	<u>EQ 5MG BASE</u>	<u>A078807 001</u>	Jun 12, 2008
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A078807 002</u>	Jun 12, 2008
<u>AB</u>	PAR PHARM INC	<u>EQ 10MG BASE</u>	<u>A070581 001</u>	Oct 17, 1985
<u>AB</u>	TEVA	<u>EQ 5MG BASE</u>	<u>A072801 001</u>	Jun 15, 1993
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A070184 001</u>	Jul 29, 1985
<u>AB</u>	VINTAGE PHARMS	<u>EQ 5MG BASE</u>	<u>A077878 001</u>	Aug 28, 2006
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A077878 002</u>	Aug 28, 2006

REGLAN

<u>AB</u>	+ ANI PHARMS	<u>EQ 5MG BASE</u>	<u>N017854 002</u>	May 05, 1987
<u>AB</u>	+!	<u>EQ 10MG BASE</u>	<u>N017854 001</u>	

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

<u>AB</u>	MYLAN	<u>2.5MG</u>	<u>A076698 001</u>	Dec 23, 2003
<u>AB</u>		<u>5MG</u>	<u>A076698 002</u>	Oct 19, 2004
<u>AB</u>		<u>10MG</u>	<u>A076698 003</u>	Oct 19, 2004
<u>AB</u>	SANDOZ	<u>2.5MG</u>	<u>A076732 001</u>	Dec 19, 2003

PRESCRIPTION DRUG PRODUCT LIST

METOLAZONE

TABLET; ORAL

METOLAZONE

<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

ZAROXOLYN

<u>AB</u>	+	LANNETT CO INC	<u>2.5MG</u>	<u>N017386</u>	<u>001</u>
<u>AB</u>	+	!	<u>5MG</u>	<u>N017386</u>	<u>002</u>
<u>AB</u>	+	!	<u>10MG</u>	<u>N017386</u>	<u>003</u>

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>		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>		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>		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>		TWI PHARMS	<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>		ZYDUS PHARMS	<u>EQ 25MG TARTRATE</u>	<u>A203894</u>	<u>001</u>	Mar 23, 2018
<u>AB</u>			<u>EQ 50MG TARTRATE</u>	<u>A203894</u>	<u>002</u>	Mar 23, 2018
<u>AB</u>			<u>EQ 100MG TARTRATE</u>	<u>A203894</u>	<u>003</u>	Mar 23, 2018
<u>AB</u>			<u>EQ 200MG TARTRATE</u>	<u>A203894</u>	<u>004</u>	Mar 23, 2018
		<u>TOPROL-XL</u>				
<u>AB</u>	+	TOPROL	<u>EQ 25MG TARTRATE</u>	<u>N019962</u>	<u>004</u>	Feb 05, 2001
<u>AB</u>	+	!	<u>EQ 50MG TARTRATE</u>	<u>N019962</u>	<u>001</u>	Jan 10, 1992
<u>AB</u>	+		<u>EQ 100MG TARTRATE</u>	<u>N019962</u>	<u>002</u>	Jan 10, 1992
<u>AB</u>	+	!	<u>EQ 200MG TARTRATE</u>	<u>N019962</u>	<u>003</u>	Jan 10, 1992

METOPROLOL TARTRATE

INJECTABLE; INJECTION

METOPROLOL TARTRATE

<u>AP</u>		BAXTER HLTHCARE CORP	<u>1MG/ML</u>	<u>A078950</u>	<u>001</u>	Apr 29, 2013
<u>AP</u>		FRESENIUS KABI USA	<u>1MG/ML</u>	<u>A091045</u>	<u>001</u>	Oct 25, 2010
<u>AP</u>		GLAND PHARMA LTD	<u>1MG/ML</u>	<u>A204205</u>	<u>001</u>	Aug 27, 2014
<u>AP</u>		HIKMA FARMACEUTICA	<u>1MG/ML</u>	<u>A077761</u>	<u>001</u>	May 30, 2007
<u>AP</u>		HOSPIRA	<u>1MG/ML</u>	<u>A074133</u>	<u>001</u>	Dec 21, 1993
<u>AP</u>			<u>1MG/ML</u>	<u>A075160</u>	<u>001</u>	Jul 06, 1998
<u>AP</u>	!		<u>1MG/ML</u>	<u>A078085</u>	<u>001</u>	Apr 29, 2008
<u>AP</u>		MYLAN ASI	<u>1MG/ML</u>	<u>A090317</u>	<u>001</u>	Apr 19, 2010
<u>AP</u>		SANDOZ INC	<u>1MG/ML</u>	<u>A077360</u>	<u>001</u>	Oct 02, 2007
<u>AP</u>		WEST-WARD PHARMS INT	<u>1MG/ML</u>	<u>A076495</u>	<u>001</u>	Jul 07, 2003

PRESCRIPTION DRUG PRODUCT LIST

METOPROLOL TARTRATE

TABLET; ORAL

LOPRESSOR

AB + US PHARMS HOLDINGS **50MG** **N017963 001**
 I
AB + **100MG** **N017963 002**

METOPROLOL TARTRATE

AB ALEMBIC PHARMS LTD **25MG** **A202871 001** May 28, 2013
AB **50MG** **A202871 002** May 28, 2013
AB **100MG** **A202871 003** May 28, 2013
AB AUROBINDO PHARMA **25MG** **A077739 001** Sep 11, 2007
AB **50MG** **A077739 002** Sep 11, 2007
AB **100MG** **A077739 003** Sep 11, 2007
AB IPCA LABS LTD **25MG** **A078459 001** Jun 17, 2008
AB **50MG** **A078459 002** Jun 17, 2008
AB **100MG** **A078459 003** Jun 17, 2008
AB MYLAN **25MG** **A076704 001** Jan 16, 2004
AB **37.5MG** **A076704 004** Mar 18, 2015
AB **50MG** **A076704 002** Jan 16, 2004
AB **75MG** **A076704 005** Mar 18, 2015
AB ! **100MG** **A076704 003** Jan 16, 2004
AB RUBICON **25MG** **A200981 001** Oct 28, 2014
AB **37.5MG** **A200981 004** Aug 21, 2019
AB **50MG** **A200981 002** Oct 28, 2014
AB **75MG** **A200981 005** Aug 21, 2019
AB **100MG** **A200981 003** Oct 28, 2014
AB SUN PHARM INDS INC **25MG** **A076670 001** Jan 15, 2004
AB **50MG** **A074644 001** Dec 10, 1996
AB **100MG** **A074644 002** Dec 10, 1996
AB YOUNGTECH PHARMS **25MG** **A208955 001** Feb 05, 2020
 INC
AB **50MG** **A208955 002** Feb 05, 2020
AB **100MG** **A208955 003** Feb 05, 2020

METRONIDAZOLE

CAPSULE; ORAL

FLAGYL

AB +! GD SEARLE LLC **375MG** **N020334 001** May 03, 1995

METRONIDAZOLE

AB ALEMBIC PHARMS LTD **375MG** **A079065 001** Jun 23, 2009
AB PAR PHARM **375MG** **A076522 001** Jan 29, 2004

CREAM; TOPICAL

METROCREAM

AB +! GALDERMA LABS LP **0.75%** **N020531 001** Sep 20, 1995

METRONIDAZOLE

AB ACP NIMBLE **0.75%** **A077549 001** Dec 19, 2007
AB FOUGERA PHARMS **0.75%** **A076408 001** May 28, 2004

NORITATE

+! VALEANT PHARMS **1%** **N020743 001** Sep 26, 1997
 NORTH

GEL; TOPICAL

METROGEL

AB +! GALDERMA LABS LP **0.75%** **N019737 001** Nov 22, 1988
AB +! **1%** **N021789 001** Jun 30, 2005

METRONIDAZOLE

AB ACP NIMBLE **0.75%** **A078178 001** Jan 19, 2011
AB FOUGERA PHARMS **0.75%** **A077018 001** Jun 06, 2006
AB TARO **0.75%** **A077819 001** Jul 18, 2006
AB **1%** **A204651 001** Mar 14, 2017
AB TOLMAR **0.75%** **A077547 001** Jul 13, 2006
AB **1%** **A090903 001** Jul 22, 2011

GEL; VAGINAL

METROGEL-VAGINAL

AB +! BAUSCH **0.75%** **N020208 001** Aug 17, 1992

METRONIDAZOLE

AB PERRIGO UK FINCO **0.75%** **A211786 001** Jul 02, 2019
AB TOLMAR **0.75%** **A077264 001** Oct 31, 2006

VANDAZOLE

BX TEVA PHARMS **0.75%** **N021806 001** May 20, 2005

NUVESSA

+! CHEMO RESEARCH SL **1.3%** **N205223 001** Mar 24, 2014

PRESCRIPTION DRUG PRODUCT LIST

METRONIDAZOLE

INJECTABLE; INJECTION

FLAGYL I.V. RTU IN PLASTIC CONTAINER

<u>AP</u>	<u>+!</u>	<u>BAXTER HLTHCARE</u>	<u>500MG/100ML</u>	<u>N018657</u>	<u>001</u>	
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METRO I.V. IN PLASTIC CONTAINER

<u>AP</u>	<u>+!</u>	<u>B BRAUN</u>	<u>500MG/100ML</u>	<u>N018900</u>	<u>001</u>	Sep 29, 1983
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METRONIDAZOLE IN PLASTIC CONTAINER

<u>AP</u>		<u>BAXTER HLTHCARE</u>	<u>500MG/100ML</u>	<u>A078084</u>	<u>001</u>	Mar 31, 2008
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<u>AP</u>	<u>+!</u>	<u>HOSPIRA</u>	<u>500MG/100ML</u>	<u>N018890</u>	<u>002</u>	Nov 18, 1983
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<u>AP</u>		<u>INFORLIFE</u>	<u>500MG/100ML</u>	<u>A206191</u>	<u>001</u>	Feb 25, 2019
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<u>AP</u>		<u>MYLAN LABS LTD</u>	<u>500MG/100ML</u>	<u>A205531</u>	<u>001</u>	May 09, 2017
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LOTION; TOPICAL

METROLOTION

<u>AB</u>	<u>+!</u>	<u>GALDERMA LABS LP</u>	<u>0.75%</u>	<u>N020901</u>	<u>001</u>	Nov 24, 1998
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METRONIDAZOLE

<u>AB</u>		<u>FOUGERA PHARMS</u>	<u>0.75%</u>	<u>A077197</u>	<u>001</u>	May 24, 2006
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TABLET; ORAL

FLAGYL

<u>AB</u>	<u>+</u>	<u>GD SEARLE LLC</u>	<u>250MG</u>	<u>N012623</u>	<u>001</u>	
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<u>AB</u>	<u>+!</u>		<u>500MG</u>	<u>N012623</u>	<u>003</u>	
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METRONIDAZOLE

<u>AB</u>		<u>ALEMBIC PHARMS LTD</u>	<u>250MG</u>	<u>A079067</u>	<u>001</u>	Mar 13, 2009
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<u>AB</u>			<u>500MG</u>	<u>A079067</u>	<u>002</u>	Mar 13, 2009
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<u>AB</u>		<u>AUROBINDO PHARMA</u>	<u>250MG</u>	<u>A203974</u>	<u>001</u>	May 29, 2015
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<u>AB</u>		<u>LTD</u>				
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<u>AB</u>			<u>500MG</u>	<u>A203974</u>	<u>002</u>	May 29, 2015
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<u>AB</u>		<u>CADILA</u>	<u>250MG</u>	<u>A206560</u>	<u>001</u>	Nov 16, 2016
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<u>AB</u>			<u>500MG</u>	<u>A206560</u>	<u>002</u>	Nov 16, 2016
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<u>AB</u>		<u>CADILA PHARMS LTD</u>	<u>250MG</u>	<u>A209794</u>	<u>001</u>	Dec 12, 2017
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<u>AB</u>			<u>500MG</u>	<u>A209794</u>	<u>002</u>	Dec 12, 2017
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<u>AB</u>		<u>FLAMINGO PHARMS</u>	<u>250MG</u>	<u>A207309</u>	<u>001</u>	May 16, 2016
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<u>AB</u>			<u>500MG</u>	<u>A207309</u>	<u>002</u>	May 16, 2016
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<u>AB</u>		<u>HERITAGE PHARMS INC</u>	<u>250MG</u>	<u>A205245</u>	<u>001</u>	Sep 23, 2015
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<u>AB</u>			<u>500MG</u>	<u>A205245</u>	<u>002</u>	Sep 23, 2015
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<u>AB</u>		<u>INNOGENIX</u>	<u>250MG</u>	<u>A070772</u>	<u>001</u>	Jul 16, 1986
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<u>AB</u>			<u>500MG</u>	<u>A070772</u>	<u>002</u>	Jul 16, 1986
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<u>AB</u>		<u>LUPIN LTD</u>	<u>250MG</u>	<u>A209096</u>	<u>001</u>	Sep 12, 2017
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<u>AB</u>			<u>500MG</u>	<u>A209096</u>	<u>002</u>	Sep 12, 2017
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<u>AB</u>		<u>ORIT LABS LLC</u>	<u>250MG</u>	<u>A208681</u>	<u>001</u>	Jun 20, 2017
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<u>AB</u>			<u>500MG</u>	<u>A208681</u>	<u>002</u>	Jun 20, 2017
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<u>AB</u>		<u>PLIVA</u>	<u>500MG</u>	<u>A070033</u>	<u>001</u>	Dec 06, 1984
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<u>AB</u>		<u>STRIDES PHARMA</u>	<u>250MG</u>	<u>A070040</u>	<u>001</u>	Jan 29, 1985
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<u>AB</u>			<u>250MG</u>	<u>A208162</u>	<u>001</u>	May 25, 2016
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<u>AB</u>			<u>500MG</u>	<u>A070039</u>	<u>001</u>	Jan 29, 1985
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<u>AB</u>			<u>500MG</u>	<u>A208162</u>	<u>002</u>	May 25, 2016
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<u>AB</u>		<u>TEVA PHARMS USA</u>	<u>250MG</u>	<u>A070027</u>	<u>001</u>	Nov 06, 1984
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<u>AB</u>		<u>UNICHEM LABS LTD</u>	<u>250MG</u>	<u>A203458</u>	<u>001</u>	Jan 22, 2014
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<u>AB</u>			<u>500MG</u>	<u>A203458</u>	<u>002</u>	Jan 22, 2014
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<u>AB</u>		<u>WATSON LABS</u>	<u>250MG</u>	<u>A070035</u>	<u>001</u>	Dec 20, 1984
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<u>AB</u>		<u>WATSON LABS INC</u>	<u>500MG</u>	<u>A070044</u>	<u>001</u>	Feb 08, 1985
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METYRAPONE

CAPSULE; ORAL

METOPIRONE

<u>+!</u>	<u>HRA PHARMA</u>	<u>250MG</u>	<u>N012911</u>	<u>002</u>	Aug 09, 1996
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METYROSINE

CAPSULE; ORAL

DEMSEER

<u>+!</u>	<u>BAUSCH</u>	<u>250MG</u>	<u>N017871</u>	<u>001</u>	
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MEXILETINE HYDROCHLORIDE

CAPSULE; ORAL

MEXILETINE HYDROCHLORIDE

	<u>TEVA</u>	<u>150MG</u>	<u>A074377</u>	<u>001</u>	May 16, 1995
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		<u>200MG</u>	<u>A074377</u>	<u>002</u>	May 16, 1995
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<u>!</u>		<u>250MG</u>	<u>A074377</u>	<u>003</u>	May 16, 1995
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PRESCRIPTION DRUG PRODUCT LIST

MICAFUNGIN SODIUM

INJECTABLE; INTRAVENOUS

MICAFUNGIN SODIUM

<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

MYCAMINE

<u>AP</u>	+! ASTELLAS	<u>EQ 50MG BASE/VIAL</u>	<u>N021506 002</u>	Mar 16, 2005
<u>AP</u>	+!	<u>EQ 100MG BASE/VIAL</u>	<u>N021506 003</u>	Jun 27, 2006

MICONAZOLE

TABLET; BUCCAL

ORAVIG

+!	FORTOVIA	50MG	N022404 001	Apr 16, 2010
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MICONAZOLE NITRATE

SUPPOSITORY; VAGINAL

MICONAZOLE NITRATE

<u>AB</u>	ACTAVIS PHARMA	<u>200MG</u>	<u>A073508 001</u>	Nov 19, 1993
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MONISTAT 3

<u>AB</u>	+! MEDTECH PRODUCTS	<u>200MG</u>	<u>N018888 001</u>	Aug 15, 1984
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MICONAZOLE NITRATE; PETROLATUM, WHITE; ZINC OXIDE

OINTMENT; TOPICAL

VUSION

+!	MYLAN	0.25%; 81.35%; 15%	N021026 001	Feb 16, 2006
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MIDAZOLAM

SPRAY; NASAL

NAYZILAM

+!	UCB INC	5MG/SPRAY	N211321 001	May 17, 2019
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MIDAZOLAM HYDROCHLORIDE

INJECTABLE; INJECTION

MIDAZOLAM HYDROCHLORIDE

<u>AP</u>	AKORN INC	<u>EQ 1MG BASE/ML</u>	<u>A075494 001</u>	Jun 30, 2000
<u>AP</u>		<u>EQ 5MG BASE/ML</u>	<u>A075494 002</u>	Jun 30, 2000
<u>AP</u>	FRESENIUS KABI USA	<u>EQ 1MG BASE/ML</u>	<u>A075154 002</u>	Jun 20, 2000
<u>AP</u>		<u>EQ 5MG BASE/ML</u>	<u>A075154 001</u>	Jun 20, 2000
<u>AP</u>	GLAND PHARMA LTD	<u>EQ 1MG BASE/ML</u>	<u>A090696 001</u>	Feb 29, 2012
<u>AP</u>		<u>EQ 5MG BASE/ML</u>	<u>A090850 001</u>	Jan 25, 2012
<u>AP</u>	! HOSPIRA	<u>EQ 1MG BASE/ML</u>	<u>A075293 001</u>	Jun 20, 2000
<u>AP</u>		<u>EQ 1MG BASE/ML</u>	<u>A075856 001</u>	Jun 13, 2002
<u>AP</u>	!	<u>EQ 5MG BASE/ML</u>	<u>A075293 002</u>	Jun 20, 2000
<u>AP</u>		<u>EQ 5MG BASE/ML</u>	<u>A075856 002</u>	Jun 13, 2002
<u>AP</u>	WEST-WARD PHARMS INT	<u>EQ 1MG BASE/ML</u>	<u>A075243 001</u>	Jun 20, 2000
<u>AP</u>		<u>EQ 1MG BASE/ML</u>	<u>A075247 002</u>	Jun 23, 2000
<u>AP</u>		<u>EQ 1MG BASE/ML</u>	<u>A075324 001</u>	Jun 20, 2000
<u>AP</u>		<u>EQ 1MG BASE/ML</u>	<u>A075421 002</u>	Jun 20, 2000
<u>AP</u>		<u>EQ 5MG BASE/ML</u>	<u>A075243 002</u>	Jun 20, 2000
<u>AP</u>		<u>EQ 5MG BASE/ML</u>	<u>A075247 001</u>	Jun 23, 2000
<u>AP</u>		<u>EQ 5MG BASE/ML</u>	<u>A075324 002</u>	Jun 20, 2000
<u>AP</u>		<u>EQ 5MG BASE/ML</u>	<u>A075421 001</u>	Jun 20, 2000

MIDAZOLAM HYDROCHLORIDE PRESERVATIVE FREE

<u>AP</u>	FRESENIUS KABI USA	<u>EQ 1MG BASE/ML</u>	<u>A203460 001</u>	Aug 22, 2014
<u>AP</u>		<u>EQ 5MG BASE/ML</u>	<u>A203460 002</u>	Aug 22, 2014
<u>AP</u>	! HOSPIRA	<u>EQ 1MG BASE/ML</u>	<u>A075857 001</u>	Jul 22, 2002
<u>AP</u>	!	<u>EQ 5MG BASE/ML</u>	<u>A075857 002</u>	Jul 22, 2002

MIDAZOLAM HYDROCHLORIDE

FRESENIUS KABI USA

EQ 5MG BASE/ML	A208878 001	Mar 28, 2017
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SOLUTION; INTRAMUSCULAR

SEIZALAM

+!	MERIDIAN MEDCL TECHN	EQ 50MG BASE/10ML (EQ 5MG BASE/ML)	N209566 001	Sep 14, 2018
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SYRUP; ORAL

MIDAZOLAM HYDROCHLORIDE

<u>AA</u>	HI TECH PHARMA	<u>EQ 2MG BASE/ML</u>	<u>A075958 001</u>	Sep 04, 2003
<u>AA</u>	! HIKMA	<u>EQ 2MG BASE/ML</u>	<u>A075873 001</u>	Apr 30, 2002
<u>AA</u>	PADDOCK LLC	<u>EQ 2MG BASE/ML</u>	<u>A076379 001</u>	May 02, 2005

PRESCRIPTION DRUG PRODUCT LISTMIDODRINE HYDROCHLORIDE

TABLET; ORAL

MIDODRINE HYDROCHLORIDE

AB	APOTEX	2.5MG	A077746 001	Sep 12, 2006
AB		5MG	A077746 002	Sep 12, 2006
AB		10MG	A077746 003	Sep 12, 2006
AB	IMPAX PHARMS	2.5MG	A076449 001	May 27, 2004
AB		5MG	A076449 002	May 27, 2004
AB		10MG	A076449 003	Dec 16, 2005
AB	MYLAN PHARMS INC	2.5MG	A076577 001	Sep 10, 2003
AB		5MG	A076577 002	Sep 10, 2003
AB		10MG	A076577 003	Sep 10, 2003
AB	PAR PHARM INC	2.5MG	A207169 001	Oct 29, 2018
AB		5MG	A207169 002	Oct 29, 2018
AB		10MG	A207169 003	Oct 29, 2018
AB	RUBICON	2.5MG	A212543 001	Aug 19, 2019
AB		5MG	A212543 002	Aug 19, 2019
AB		10MG	A212543 003	Aug 19, 2019
AB	UNIQUE PHARM LABS	2.5MG	A207613 001	Nov 02, 2018
AB		5MG	A207613 002	Nov 02, 2018
AB		10MG	A207613 003	Nov 02, 2018
ORVATEN				
AB	UPSHER SMITH LABS	2.5MG	A076725 001	Nov 03, 2004
AB	!	5MG	A076725 002	Nov 03, 2004
AB		10MG	A076725 003	Nov 03, 2004

MIDOSTAURIN

CAPSULE; ORAL

RYDAPT

+! NOVARTIS 25MG N207997 001 Apr 28, 2017

MIFEPRISTONE

TABLET; ORAL

MIFEPREX**AB** +! DANCO LABS LLC **200MG** **N020687 001** Sep 28, 2000MIFEPRISTONE**AB** GENBIOPRO **200MG** **A091178 001** Apr 11, 2019

KORLYM

+! CORCEPT THERAP 300MG N202107 001 Feb 17, 2012

MIGALASTAT HYDROCHLORIDE

CAPSULE; ORAL

GALAFOLD

+! AMICUS THERAPS US EQ 123MG BASE N208623 001 Aug 10, 2018

MIGLITOL

TABLET; ORAL

GLYSET**AB** +! PHARMACIA AND UPJOHN **25MG** **N020682 001** Dec 18, 1996**AB** + **50MG** **N020682 002** Dec 18, 1996**AB** + **100MG** **N020682 003** Dec 18, 1996MIGLITOL**AB** ORIENT PHARMA CO LTD **25MG** **A203965 001** Feb 24, 2015**AB** **50MG** **A203965 002** Feb 24, 2015**AB** **100MG** **A203965 003** Feb 24, 2015MIGLUSTAT

CAPSULE; ORAL

MIGLUSTAT**AB** AMERIGEN PHARMS LTD **100MG** **A208342 001** Apr 17, 2018ZAVESCA**AB** +! ACTELION PHARMS LTD **100MG** **N021348 001** Jul 31, 2003MILNACIPRAN HYDROCHLORIDE

TABLET; ORAL

SAVELLA

+ ALLERGAN 12.5MG N022256 001 Jan 14, 2009

+ 25MG N022256 002 Jan 14, 2009

+! 50MG N022256 003 Jan 14, 2009

+ 100MG N022256 004 Jan 14, 2009

PRESCRIPTION DRUG PRODUCT LIST

MILRINONE LACTATE

INJECTABLE; INJECTION

MILRINONE LACTATE

<u>AP</u>	FRESENIUS KABI USA	<u>EQ 1MG BASE/ML</u>	<u>A075936 001</u>	May 28, 2002
<u>AP</u>	! HIKMA FARMACEUTICA	<u>EQ 1MG BASE/ML</u>	<u>A077966 001</u>	Dec 03, 2010
<u>AP</u>	HOSPIRA INC	<u>EQ 1MG BASE/ML</u>	<u>A203280 001</u>	Sep 03, 2014
<u>AP</u>	INTL MEDICATED	<u>EQ 1MG BASE/ML</u>	<u>A076013 001</u>	Aug 02, 2002
<u>AP</u>	WEST-WARD PHARMS INT	<u>EQ 1MG BASE/ML</u>	<u>A075530 001</u>	May 28, 2002
<u>AP</u>		<u>EQ 1MG BASE/ML</u>	<u>A075660 001</u>	May 28, 2002
	<u>MILRINONE LACTATE IN DEXTROSE 5%</u>			
<u>AP</u>	WOODWARD	<u>EQ 40MG BASE/200ML (EQ 0.2MG BASE/ML)</u>	<u>A077151 002</u>	Jul 20, 2005
	<u>MILRINONE LACTATE IN DEXTROSE 5% IN PLASTIC CONTAINER</u>			
<u>AP</u>	! BAXTER HLTHCARE	<u>EQ 20MG BASE/100ML (EQ 0.2MG BASE/ML)</u>	<u>A075834 001</u>	May 28, 2002
<u>AP</u>	! HOSPIRA	<u>EQ 40MG BASE/200ML (EQ 0.2MG BASE/ML)</u>	<u>A075834 002</u>	May 28, 2002
<u>AP</u>	HOSPIRA	<u>EQ 20MG BASE/100ML (EQ 0.2MG BASE/ML)</u>	<u>A075885 001</u>	May 28, 2002
<u>AP</u>		<u>EQ 40MG BASE/200ML (EQ 0.2MG BASE/ML)</u>	<u>A075885 002</u>	May 28, 2002
<u>AP</u>	WEST-WARD PHARMS INT	<u>EQ 20MG BASE/100ML (EQ 0.2MG BASE/ML)</u>	<u>A078113 001</u>	May 21, 2008
<u>AP</u>		<u>EQ 40MG BASE/200ML (EQ 0.2MG BASE/ML)</u>	<u>A078113 002</u>	May 21, 2008
	<u>MILRINONE LACTATE IN PLASTIC CONTAINER</u>			
<u>AP</u>	HIKMA FARMACEUTICA	<u>EQ 20MG BASE/100ML (EQ 0.2MG BASE/ML)</u>	<u>A090038 001</u>	Jan 21, 2010
<u>AP</u>		<u>EQ 40MG BASE/200ML (EQ 0.2MG BASE/ML)</u>	<u>A090038 002</u>	Jan 21, 2010

MILTEFOSINE

CAPSULE; ORAL

IMPAVIDO

+! KNIGHT THERAPS

50MG

N204684 001 Mar 19, 2014

MINOCYCLINE HYDROCHLORIDE

AEROSOL, FOAM; TOPICAL

AMZEEQ

+! FOAMIX

EQ 4% BASE

N212379 001 Oct 18, 2019

CAPSULE; ORAL

DYNACIN

<u>AB</u>	CNTY LINE PHARMS	<u>EQ 50MG BASE</u>	<u>A063067 003</u>	Aug 14, 1990
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A063067 002</u>	Sep 15, 1999
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A063067 001</u>	Jul 31, 1990

MINOCIN

<u>AB</u>	+ BAUSCH	<u>EQ 50MG BASE</u>	<u>N050649 001</u>	May 31, 1990
<u>AB</u>	+ BAUSCH	<u>EQ 100MG BASE</u>	<u>N050649 002</u>	May 31, 1990

MINOCYCLINE HYDROCHLORIDE

<u>AB</u>	AUROBINDO PHARMA	<u>EQ 50MG BASE</u>	<u>A065470 001</u>	Mar 11, 2008
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A065470 002</u>	Mar 11, 2008
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A065470 003</u>	Mar 11, 2008
<u>AB</u>	IMPAX LABS	<u>EQ 50MG BASE</u>	<u>A065005 001</u>	Mar 23, 1999
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A065005 003</u>	Apr 18, 2001
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A065005 002</u>	Mar 23, 1999
<u>AB</u>	SUN PHARM INDS INC	<u>EQ 50MG BASE</u>	<u>A090867 001</u>	May 13, 2013
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A090867 002</u>	May 13, 2013
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A090867 003</u>	May 13, 2013
<u>AB</u>	TORRENT	<u>EQ 50MG BASE</u>	<u>A065062 001</u>	Nov 30, 2000
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A065062 002</u>	Nov 30, 2000
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A065062 003</u>	Nov 30, 2000
<u>AB</u>	WATSON LABS	<u>EQ 75MG BASE</u>	<u>A063065 002</u>	Jun 10, 1999
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A063065 001</u>	Dec 30, 1991
<u>AB</u>	WATSON LABS TEVA	<u>EQ 50MG BASE</u>	<u>A063181 001</u>	Dec 30, 1991
<u>AB</u>	ZYDUS	<u>EQ 50MG BASE</u>	<u>A063011 001</u>	Mar 02, 1992
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A063009 002</u>	Aug 12, 2003
<u>AB</u>	!	<u>EQ 100MG BASE</u>	<u>A063009 001</u>	Mar 02, 1992

CAPSULE, EXTENDED RELEASE; ORAL

XIMINO

JOURNEY

EQ 45MG BASE

N201922 001 Jul 11, 2012

EQ 90MG BASE

N201922 003 Jul 11, 2012

EQ 135MG BASE

N201922 005 Jul 11, 2012

INJECTABLE; INJECTION

MINOCIN

+! REMPEX PHARMS

EQ 100MG BASE/VIAL

N050444 001

POWDER, EXTENDED RELEASE; DENTAL

ARESTIN

+! ORAPHARMA

EQ 1MG BASE

N050781 001 Feb 16, 2001

PRESCRIPTION DRUG PRODUCT LIST

3-302 (of 453)

MINOCYCLINE HYDROCHLORIDE

TABLET; ORAL

MINOCYCLINE HYDROCHLORIDE

<u>AB</u>	DR REDDYS LABS LTD	<u>EQ 50MG BASE</u>	<u>A065436 001</u>	Dec 26, 2007
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A065436 002</u>	Dec 26, 2007
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A065436 003</u>	Dec 26, 2007
<u>AB</u>	PAR PHARM	<u>EQ 50MG BASE</u>	<u>A065131 001</u>	Apr 16, 2003
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A065131 002</u>	Apr 16, 2003
<u>AB</u>	!	<u>EQ 100MG BASE</u>	<u>A065131 003</u>	Apr 16, 2003
<u>AB</u>	SUN PHARM INDUSTRIES	<u>EQ 50MG BASE</u>	<u>A090217 001</u>	Jan 29, 2016
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A090217 002</u>	Jan 29, 2016
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A090217 003</u>	Jan 29, 2016
<u>AB</u>	TORRENT	<u>EQ 50MG BASE</u>	<u>A065156 001</u>	Jan 06, 2004
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A065156 002</u>	Jan 06, 2004
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A065156 003</u>	Jan 06, 2004

TABLET, EXTENDED RELEASE; ORAL

MINOCYCLINE HYDROCHLORIDE

<u>AB</u>	ALKEM LABS LTD	<u>EQ 45MG BASE</u>	<u>A204453 001</u>	Sep 28, 2016
<u>AB</u>		<u>EQ 55MG BASE</u>	<u>A204453 008</u>	Dec 19, 2019
<u>AB</u>		<u>EQ 65MG BASE</u>	<u>A204453 006</u>	Mar 16, 2018
<u>AB</u>		<u>EQ 80MG BASE</u>	<u>A204453 002</u>	Sep 28, 2016
<u>AB</u>		<u>EQ 90MG BASE</u>	<u>A204453 003</u>	Sep 28, 2016
<u>AB</u>		<u>EQ 105MG BASE</u>	<u>A204453 004</u>	Sep 28, 2016
<u>AB</u>		<u>EQ 115MG BASE</u>	<u>A204453 007</u>	Mar 16, 2018
<u>AB</u>		<u>EQ 135MG BASE</u>	<u>A204453 005</u>	Sep 28, 2016
<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>	BARR LABS INC	<u>EQ 65MG BASE</u>	<u>A065485 004</u>	May 18, 2012
<u>AB</u>		<u>EQ 80MG BASE</u>	<u>A065485 007</u>	Apr 26, 2017
<u>AB</u>		<u>EQ 105MG BASE</u>	<u>A065485 008</u>	Apr 26, 2017
<u>AB</u>		<u>EQ 115MG BASE</u>	<u>A065485 005</u>	May 18, 2012
<u>AB</u>	LUPIN LTD	<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>	MYLAN	<u>EQ 80MG BASE</u>	<u>A203443 002</u>	Aug 22, 2014
<u>AB</u>		<u>EQ 105MG BASE</u>	<u>A203443 003</u>	Aug 22, 2014
<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>	SIDMAK LABS INDIA	<u>EQ 45MG BASE</u>	<u>A204394 001</u>	Dec 30, 2015
<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 135MG BASE</u>	<u>A204394 007</u>	Dec 30, 2015
<u>AB</u>	SUN PHARM INDS LTD	<u>EQ 45MG BASE</u>	<u>A091118 001</u>	Sep 25, 2014
<u>AB</u>		<u>EQ 65MG BASE</u>	<u>A091118 003</u>	Dec 03, 2019
<u>AB</u>		<u>EQ 80MG BASE</u>	<u>A091118 004</u>	Sep 25, 2014
<u>AB</u>		<u>EQ 90MG BASE</u>	<u>A091118 005</u>	Sep 25, 2014
<u>AB</u>		<u>EQ 105MG BASE</u>	<u>A091118 006</u>	Sep 25, 2014
<u>AB</u>		<u>EQ 115MG BASE</u>	<u>A091118 007</u>	Dec 03, 2019
<u>AB</u>		<u>EQ 135MG BASE</u>	<u>A091118 008</u>	Sep 25, 2014
<u>AB</u>	ZYDUS PHARMS	<u>EQ 45MG BASE</u>	<u>A203553 001</u>	Nov 16, 2017
<u>AB</u>		<u>EQ 80MG BASE</u>	<u>A203553 004</u>	Nov 16, 2017
<u>AB</u>		<u>EQ 90MG BASE</u>	<u>A203553 005</u>	Nov 16, 2017
<u>AB</u>		<u>EQ 105MG BASE</u>	<u>A203553 006</u>	Nov 16, 2017
<u>AB</u>		<u>EQ 135MG BASE</u>	<u>A203553 008</u>	Nov 16, 2017
<u>SOLODYN</u>				
<u>AB</u>	+ MEDICIS	<u>EQ 55MG BASE</u>	<u>N050808 008</u>	Aug 27, 2010
<u>AB</u>	+	<u>EQ 65MG BASE</u>	<u>N050808 004</u>	Jul 23, 2009
<u>AB</u>	+	<u>EQ 80MG BASE</u>	<u>N050808 007</u>	Aug 27, 2010
<u>AB</u>	+	<u>EQ 105MG BASE</u>	<u>N050808 006</u>	Aug 27, 2010
<u>AB</u>	+	<u>EQ 115MG BASE</u>	<u>N050808 005</u>	Jul 23, 2009
<u>MINOLIRA</u>				
	EPI HLTH	EQ 105MG BASE	N209269 001	May 08, 2017
		EQ 135MG BASE	N209269 002	May 08, 2017

PRESCRIPTION DRUG PRODUCT LIST

MINOXIDIL

TABLET; ORAL

MINOXIDIL

<u>AB</u>	PAR PHARM	<u>2.5MG</u>	<u>A071826</u>	<u>001</u>	Nov 14, 1988
<u>AB</u>		<u>10MG</u>	<u>A071839</u>	<u>001</u>	Nov 14, 1988
<u>AB</u>	SUN PHARM INDUSTRIES	<u>2.5MG</u>	<u>A072709</u>	<u>002</u>	Dec 14, 1995
<u>AB</u>		<u>10MG</u>	<u>A072709</u>	<u>001</u>	Dec 14, 1995
<u>AB</u>	WATSON LABS	<u>2.5MG</u>	<u>A071344</u>	<u>001</u>	Mar 03, 1987
<u>AB</u>	!	<u>10MG</u>	<u>A071345</u>	<u>001</u>	Mar 03, 1987

MIRABEGRON

TABLET, EXTENDED RELEASE; ORAL

MYRBETRIQ

+	!	APGDI	25MG	N202611	001	Jun 28, 2012
+	!		50MG	N202611	002	Jun 28, 2012

MIRTAZAPINE

TABLET; ORAL

MIRTAZAPINE

<u>AB</u>	APOTEX INC	<u>15MG</u>	<u>A077666</u>	<u>001</u>	Aug 22, 2007
<u>AB</u>		<u>30MG</u>	<u>A077666</u>	<u>002</u>	Aug 22, 2007
<u>AB</u>		<u>45MG</u>	<u>A077666</u>	<u>003</u>	Aug 22, 2007
<u>AB</u>	AUROBINDO	<u>7.5MG</u>	<u>A076921</u>	<u>001</u>	Oct 22, 2004
<u>AB</u>		<u>15MG</u>	<u>A076921</u>	<u>002</u>	Oct 22, 2004
<u>AB</u>		<u>30MG</u>	<u>A076921</u>	<u>003</u>	Oct 22, 2004
<u>AB</u>		<u>45MG</u>	<u>A076921</u>	<u>004</u>	Oct 22, 2004
<u>AB</u>	MYLAN	<u>15MG</u>	<u>A076122</u>	<u>001</u>	Jun 19, 2003
<u>AB</u>		<u>30MG</u>	<u>A076122</u>	<u>002</u>	Jun 19, 2003
<u>AB</u>		<u>45MG</u>	<u>A076122</u>	<u>003</u>	Jun 19, 2003
<u>AB</u>	SUN PHARM INDS INC	<u>7.5MG</u>	<u>A076541</u>	<u>004</u>	Apr 22, 2004
<u>AB</u>		<u>15MG</u>	<u>A076541</u>	<u>001</u>	Apr 22, 2004
<u>AB</u>		<u>30MG</u>	<u>A076541</u>	<u>002</u>	Apr 22, 2004
<u>AB</u>		<u>45MG</u>	<u>A076541</u>	<u>003</u>	Apr 22, 2004
<u>AB</u>	TEVA	<u>15MG</u>	<u>A076119</u>	<u>001</u>	Jan 24, 2003
<u>AB</u>		<u>30MG</u>	<u>A076119</u>	<u>002</u>	Jan 24, 2003
<u>AB</u>		<u>45MG</u>	<u>A076119</u>	<u>003</u>	Jun 19, 2003
<u>AB</u>	UPSHER SMITH LABS	<u>15MG</u>	<u>A076219</u>	<u>001</u>	Jun 19, 2003
<u>AB</u>		<u>30MG</u>	<u>A076219</u>	<u>002</u>	Jun 19, 2003
<u>AB</u>		<u>45MG</u>	<u>A076219</u>	<u>003</u>	Jun 19, 2003

REMERON

<u>AB</u>	+	!	ORGANON USA INC	<u>15MG</u>	<u>N020415</u>	<u>001</u>	Jun 14, 1996
<u>AB</u>	+			<u>30MG</u>	<u>N020415</u>	<u>002</u>	Jun 14, 1996

TABLET, ORALLY DISINTEGRATING; ORAL

MIRTAZAPINE

<u>AB</u>	AUROBINDO PHARMA LTD	<u>15MG</u>	<u>A077376</u>	<u>002</u>	Dec 08, 2005
<u>AB</u>		<u>30MG</u>	<u>A077376</u>	<u>003</u>	Dec 08, 2005
<u>AB</u>		<u>45MG</u>	<u>A077376</u>	<u>004</u>	Feb 28, 2006
<u>AB</u>	IMPAX LABS INC	<u>15MG</u>	<u>A076901</u>	<u>001</u>	Jun 28, 2005
<u>AB</u>		<u>30MG</u>	<u>A076901</u>	<u>002</u>	Jun 28, 2005
<u>AB</u>		<u>45MG</u>	<u>A076901</u>	<u>003</u>	Jun 28, 2005
<u>AB</u>	ZYDUS PHARMS	<u>15MG</u>	<u>A205798</u>	<u>001</u>	Jun 01, 2017
<u>AB</u>		<u>30MG</u>	<u>A205798</u>	<u>002</u>	Jun 01, 2017
<u>AB</u>		<u>45MG</u>	<u>A205798</u>	<u>003</u>	Jun 01, 2017

REMERON SOLTAB

<u>AB</u>	+	!	ORGANON USA INC	<u>15MG</u>	<u>N021208</u>	<u>001</u>	Jan 12, 2001
<u>AB</u>	+			<u>30MG</u>	<u>N021208</u>	<u>002</u>	Jan 12, 2001
<u>AB</u>	+			<u>45MG</u>	<u>N021208</u>	<u>003</u>	Jan 12, 2001

MISOPROSTOL

TABLET; ORAL

CYTOTEC

<u>AB</u>	+		GD SEARLE LLC	<u>0.1MG</u>	<u>N019268</u>	<u>003</u>	Sep 21, 1990
<u>AB</u>	+	!		<u>0.2MG</u>	<u>N019268</u>	<u>001</u>	Dec 27, 1988

MISOPROSTOL

<u>AB</u>			NOVEL LABS INC	<u>0.1MG</u>	<u>A091667</u>	<u>001</u>	Jul 25, 2012
<u>AB</u>				<u>0.2MG</u>	<u>A091667</u>	<u>002</u>	Jul 25, 2012

PRESCRIPTION DRUG PRODUCT LIST

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</u>	<u>001</u>	Apr 30, 1998
<u>AP</u>	!		<u>20MG/VIAL</u>	<u>A064144</u>	<u>002</u>	Apr 30, 1998
<u>AP</u>	!		<u>40MG/VIAL</u>	<u>A064144</u>	<u>003</u>	Aug 11, 2009
<u>AP</u>		MYLAN LABS LTD	<u>5MG/VIAL</u>	<u>A202670</u>	<u>001</u>	Oct 13, 2017
<u>AP</u>			<u>20MG/VIAL</u>	<u>A202670</u>	<u>002</u>	Oct 13, 2017
<u>AP</u>			<u>40MG/VIAL</u>	<u>A203386</u>	<u>001</u>	Oct 13, 2017
<u>AP</u>		WEST-WARD PHARMS INT	<u>5MG/VIAL</u>	<u>A064180</u>	<u>001</u>	Dec 23, 1999
<u>AP</u>			<u>20MG/VIAL</u>	<u>A064117</u>	<u>002</u>	Apr 19, 1995
<u>AP</u>			<u>20MG/VIAL</u>	<u>A064180</u>	<u>002</u>	Dec 23, 1999
<u>AP</u>			<u>40MG/VIAL</u>	<u>A064117</u>	<u>003</u>	Jun 02, 1999

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</u>	<u>001</u>	Apr 11, 2006
<u>AP</u>			<u>EQ 25MG BASE/12.5ML (EQ 2MG BASE/ML)</u>	<u>A077496</u>	<u>002</u>	Apr 11, 2006
<u>AP</u>			<u>EQ 30MG BASE/15ML (EQ 2MG BASE/ML)</u>	<u>A077496</u>	<u>003</u>	Apr 11, 2006
<u>AP</u>	!	HOSPIRA	<u>EQ 20MG BASE/10ML (EQ 2MG BASE/ML)</u>	<u>A076871</u>	<u>001</u>	Apr 11, 2006
<u>AP</u>	!		<u>EQ 25MG BASE/12.5ML (EQ 2MG BASE/ML)</u>	<u>A076871</u>	<u>002</u>	Apr 11, 2006
<u>AP</u>	!		<u>EQ 30MG BASE/15ML (EQ 2MG BASE/ML)</u>	<u>A076871</u>	<u>003</u>	Apr 11, 2006
<u>AP</u>		TEVA PHARMS USA	<u>EQ 20MG BASE/10ML (EQ 2MG BASE/ML)</u>	<u>A077356</u>	<u>001</u>	Apr 11, 2006
<u>AP</u>			<u>EQ 25MG BASE/12.5ML (EQ 2MG BASE/ML)</u>	<u>A077356</u>	<u>002</u>	Apr 11, 2006
<u>AP</u>			<u>EQ 30MG BASE/15ML (EQ 2MG BASE/ML)</u>	<u>A077356</u>	<u>003</u>	Apr 11, 2006
<u>AP</u>		WEST-WARD PHARMS INT	<u>EQ 20MG BASE/10ML (EQ 2MG BASE/ML)</u>	<u>A076611</u>	<u>001</u>	Apr 11, 2006
<u>AP</u>			<u>EQ 25MG BASE/12.5ML (EQ 2MG BASE/ML)</u>	<u>A076611</u>	<u>002</u>	Apr 11, 2006
<u>AP</u>			<u>EQ 30MG BASE/15ML (EQ 2MG BASE/ML)</u>	<u>A076611</u>	<u>003</u>	Apr 11, 2006

MODAFINIL

TABLET; ORAL

MODAFINIL

<u>AB</u>		ALEMBC PHARMS LTD	<u>100MG</u>	<u>A202700</u>	<u>001</u>	Oct 18, 2012
<u>AB</u>			<u>200MG</u>	<u>A202700</u>	<u>002</u>	Oct 18, 2012
<u>AB</u>		APOTEX INC	<u>100MG</u>	<u>A077667</u>	<u>001</u>	Feb 03, 2014
<u>AB</u>			<u>200MG</u>	<u>A077667</u>	<u>002</u>	Feb 03, 2014
<u>AB</u>		APPCO	<u>100MG</u>	<u>A207196</u>	<u>001</u>	Aug 16, 2017
<u>AB</u>			<u>200MG</u>	<u>A207196</u>	<u>002</u>	Aug 16, 2017
<u>AB</u>		AUROBINDO PHARMA LTD	<u>100MG</u>	<u>A202566</u>	<u>001</u>	Sep 27, 2012
<u>AB</u>			<u>200MG</u>	<u>A202566</u>	<u>002</u>	Sep 27, 2012
<u>AB</u>		CADILA	<u>100MG</u>	<u>A209966</u>	<u>001</u>	Sep 14, 2017
<u>AB</u>			<u>200MG</u>	<u>A209966</u>	<u>002</u>	Sep 14, 2017
<u>AB</u>		MYLAN PHARMS INC	<u>100MG</u>	<u>A076594</u>	<u>001</u>	Jul 16, 2012
<u>AB</u>			<u>200MG</u>	<u>A076594</u>	<u>002</u>	Jul 16, 2012
<u>AB</u>		ORCHID HLTHCARE	<u>100MG</u>	<u>A078963</u>	<u>001</u>	Sep 26, 2012
<u>AB</u>			<u>200MG</u>	<u>A078963</u>	<u>002</u>	Sep 26, 2012
<u>AB</u>		WATSON LABS INC	<u>100MG</u>	<u>A076715</u>	<u>001</u>	Nov 01, 2012
<u>AB</u>			<u>200MG</u>	<u>A076715</u>	<u>002</u>	Nov 01, 2012

PROVIGIL

<u>AB</u>	+	CEPHALON	<u>100MG</u>	<u>N020717</u>	<u>001</u>	Dec 24, 1998
<u>AB</u>	+		<u>200MG</u>	<u>N020717</u>	<u>002</u>	Dec 24, 1998

MOEXIPRIL HYDROCHLORIDE

TABLET; ORAL

MOEXIPRIL HYDROCHLORIDE

<u>AB</u>		APOTEX INC	<u>7.5MG</u>	<u>A078454</u>	<u>001</u>	Jun 02, 2008
<u>AB</u>			<u>15MG</u>	<u>A078454</u>	<u>002</u>	Jun 02, 2008
<u>AB</u>		CHARTWELL RX	<u>7.5MG</u>	<u>A077536</u>	<u>001</u>	Nov 30, 2006
<u>AB</u>			<u>15MG</u>	<u>A077536</u>	<u>002</u>	Nov 30, 2006
<u>AB</u>		GLENMARK GENERICS	<u>7.5MG</u>	<u>A090416</u>	<u>001</u>	Mar 30, 2010
<u>AB</u>			<u>15MG</u>	<u>A090416</u>	<u>002</u>	Mar 30, 2010
<u>AB</u>		TEVA	<u>7.5MG</u>	<u>A076204</u>	<u>001</u>	May 08, 2003
<u>AB</u>	!		<u>15MG</u>	<u>A076204</u>	<u>002</u>	May 08, 2003

PRESCRIPTION DRUG PRODUCT LIST

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

MOMETASONE FUROATE

AEROSOL, METERED; INHALATION

ASMANEX HFA

+ MERCK SHARP DOHME

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** ACP NIMBLE0.1%A077447 001 May 22, 2006**AB** ANDA REPOSITORY0.1%A076591 001 Apr 18, 2007**AB** FOUGERA PHARMS0.1%A076171 001 Apr 08, 2005**AB** ! GLENMARK GENERICS0.1%A078541 001 May 28, 2008**AB** TARO0.1%A076679 001 Dec 21, 2004

IMPLANT; IMPLANTATION

SINUVA

+ INTERSECT ENT INC

1.35MG

N209310 001 Dec 08, 2017

LOTION; TOPICAL

ELOCON**AB** +! MERCK SHARP DOHME0.1%N019796 001 Mar 30, 1989MOMETASONE FUROATE**AB** ACP NIMBLE0.1%A077678 001 Nov 21, 2007**AB** ANDA REPOSITORY0.1%A076499 001 Nov 21, 2007**AB** FOUGERA PHARMS0.1%A075919 001 Nov 29, 2007**AB** GLENMARK GENERICS0.1%A090506 001 Aug 09, 2010**AB** PERRIGO ISRAEL0.1%A077180 001 Apr 06, 2005**AB** TARO0.1%A076788 001 Mar 15, 2006

OINTMENT; TOPICAL

MOMETASONE FUROATE**AB** ACP NIMBLE0.1%A077401 001 Jun 20, 2006**AB** ANDA REPOSITORY0.1%A076481 001 Nov 14, 2003**AB** FOUGERA PHARMS0.1%A077061 001 Mar 28, 2005**AB** ! GLENMARK GENERICS0.1%A078571 001 May 28, 2008**AB** PERRIGO NEW YORK0.1%A076067 001 Mar 18, 2002**AB** TORRENT0.1%A207899 001 Jul 13, 2018

POWDER; INHALATION

ASMANEX TWISTHALER

+ MERCK SHARP DOHME

0.11MG/INH

N021067 002 Feb 01, 2008

+!

0.22MG/INH

N021067 001 Mar 30, 2005

SPRAY, METERED; NASAL

MOMETASONE FUROATE**AB** AMNEAL PHARMSEQ 0.05MG BASE/SPRAYA207989 001 Apr 03, 2017**AB** APOTEX INCEQ 0.05MG BASE/SPRAYA091161 001 Mar 22, 2016NASONEX**AB** +! MERCK SHARP DOHMEEQ 0.05MG BASE/SPRAYN020762 001 Oct 01, 1997MONTELUKAST SODIUM

GRANULE; ORAL

MONTELUKAST SODIUM**AB** AJANTA PHARMA LTDEQ 4MG BASE/PACKETA203438 001 Jul 31, 2015**AB** AUROBINDO PHARMA LTDEQ 4MG BASE/PACKETA213471 001 Feb 18, 2020**AB** DR REDDYS LABS LTDEQ 4MG BASE/PACKETA202906 001 Sep 17, 2012**AB** TEVA PHARMSEQ 4MG BASE/PACKETA090955 001 Aug 03, 2012**AB** TORRENTEQ 4MG BASE/PACKETA210431 001 Jul 31, 2018SINGULAIR**AB** +! MERCKEQ 4MG BASE/PACKETN021409 001 Jul 26, 2002

TABLET; ORAL

MONTELUKAST SODIUM**AB** ACCORD HLTHCAREEQ 10MG BASEA202717 001 Sep 21, 2012**AB** AJANTA PHARMA LTDEQ 10MG BASEA203432 001 Jul 31, 2015**AB** AMNEAL PHARMSEQ 10MG BASEA204604 001 Sep 04, 2015**AB** ANBISON LABEQ 10MG BASEA205683 001 Jan 12, 2016**AB** AUROBINDO PHARMA LTDEQ 10MG BASEA202468 001 Aug 03, 2012**AB** CIPLAEQ 10MG BASEA207463 001 Oct 28, 2016**AB** CSPC OUYIEQ 10MG BASEA209012 001 Apr 24, 2017**AB** DR REDDYS LABS LTDEQ 10MG BASEA201582 001 Aug 06, 2012

PRESCRIPTION DRUG PRODUCT LIST

MONTELUKAST SODIUM

TABLET; ORAL

MONTELUKAST SODIUM

<u>AB</u>	GLENMARK GENERICS	<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>	L PERRIGO CO	<u>EQ 10MG BASE</u>	<u>A206112 001</u>	Apr 26, 2017
<u>AB</u>	LANNETT CO INC	<u>EQ 10MG BASE</u>	<u>A201522 001</u>	Aug 03, 2012
<u>AB</u>	MACLEODS PHARMS LTD	<u>EQ 10MG BASE</u>	<u>A203366 001</u>	Sep 11, 2014
<u>AB</u>	SANDOZ INC	<u>EQ 10MG BASE</u>	<u>A200889 001</u>	Aug 03, 2012
<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 LABS LTD	<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> ! MSD MERCK CO	<u>EQ 10MG BASE</u>	<u>N020829 002</u>	Feb 20, 1998
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TABLET, CHEWABLE; ORAL

MONTELUKAST SODIUM

<u>AB</u>	AJANTA PHARMA LTD	<u>EQ 4MG BASE</u>	<u>A203328 001</u>	Jul 31, 2015
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A203328 002</u>	Jul 31, 2015
<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>	CIPLA	<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>	CSPC OUYI	<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>	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>	HIKMA	<u>EQ 4MG BASE</u>	<u>A091128 001</u>	Aug 03, 2012
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A091128 002</u>	Aug 03, 2012
<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>	SANDOZ INC	<u>EQ 4MG BASE</u>	<u>A091414 001</u>	Aug 03, 2012
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A091414 002</u>	Aug 03, 2012
<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>	UNICHEM LABS LTD	<u>EQ 4MG BASE</u>	<u>A208621 001</u>	Jul 02, 2018
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A208621 002</u>	Jul 02, 2018
<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> MSD MERCK CO	<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

KADIAN

<u>AB</u>	<u>+</u> ! ALLERGAN	<u>10MG</u>	<u>N020616 008</u>	Apr 20, 2007
<u>AB</u>	<u>+</u>	<u>20MG</u>	<u>N020616 001</u>	Jul 03, 1996
<u>AB</u>	<u>+</u>	<u>30MG</u>	<u>N020616 004</u>	Mar 09, 2001
<u>AB</u>	<u>+</u>	<u>40MG</u>	<u>N020616 009</u>	Jul 09, 2012
<u>AB</u>	<u>+</u>	<u>50MG</u>	<u>N020616 002</u>	Jul 03, 1996
<u>AB</u>	<u>+</u>	<u>60MG</u>	<u>N020616 005</u>	Mar 09, 2001
<u>AB</u>	<u>+</u>	<u>70MG</u>	<u>N020616 010</u>	Jul 09, 2012
<u>AB</u>	<u>+</u>	<u>80MG</u>	<u>N020616 006</u>	Oct 27, 2006
<u>AB</u>	<u>+</u> !	<u>100MG</u>	<u>N020616 003</u>	Jul 03, 1996

MORPHINE SULFATE

<u>AB</u>	IMPAX LABS INC	<u>20MG</u>	<u>A200411 001</u>	Apr 12, 2016
<u>AB</u>		<u>30MG</u>	<u>A200411 002</u>	Apr 12, 2016
<u>AB</u>		<u>40MG</u>	<u>A200411 007</u>	Jul 25, 2018
<u>AB</u>		<u>50MG</u>	<u>A200411 003</u>	Apr 12, 2016
<u>AB</u>		<u>60MG</u>	<u>A200411 004</u>	Apr 12, 2016
<u>AB</u>		<u>80MG</u>	<u>A200411 005</u>	Apr 12, 2016

PRESCRIPTION DRUG PRODUCT LIST

MORPHINE SULFATE

CAPSULE, EXTENDED RELEASE; ORAL

MORPHINE SULFATE

<u>AB</u>		<u>100MG</u>	<u>A200411 006</u>	Apr 12, 2016
<u>AB</u>	PAR PHARM INC	<u>20MG</u>	<u>A200812 001</u>	Nov 10, 2011
<u>AB</u>		<u>30MG</u>	<u>A200812 002</u>	Nov 10, 2011
<u>AB</u>		<u>50MG</u>	<u>A200812 003</u>	Nov 10, 2011
<u>AB</u>		<u>60MG</u>	<u>A200812 004</u>	Nov 10, 2011
<u>AB</u>		<u>80MG</u>	<u>A200812 005</u>	Nov 10, 2011
<u>AB</u>		<u>100MG</u>	<u>A200812 006</u>	Nov 10, 2011
<u>AB</u>	TEVA PHARMS USA	<u>20MG</u>	<u>A202718 001</u>	Dec 29, 2014
<u>AB</u>		<u>30MG</u>	<u>A202718 002</u>	Dec 29, 2014
<u>AB</u>		<u>40MG</u>	<u>A202718 007</u>	Jun 03, 2015
<u>AB</u>		<u>50MG</u>	<u>A202718 003</u>	Dec 29, 2014
<u>AB</u>		<u>60MG</u>	<u>A202718 004</u>	Dec 29, 2014
<u>AB</u>		<u>70MG</u>	<u>A202718 008</u>	Jun 03, 2015
<u>AB</u>		<u>80MG</u>	<u>A202718 005</u>	Dec 29, 2014
<u>AB</u>		<u>100MG</u>	<u>A202718 006</u>	Dec 29, 2014
<u>AB</u>	UPSHER SMITH LABS	<u>10MG</u>	<u>A202104 001</u>	Jun 03, 2013
<u>AB</u>		<u>20MG</u>	<u>A202104 002</u>	Jun 03, 2013
<u>AB</u>		<u>30MG</u>	<u>A202104 003</u>	Jun 03, 2013
<u>AB</u>		<u>50MG</u>	<u>A202104 004</u>	Jun 03, 2013
<u>AB</u>		<u>60MG</u>	<u>A202104 005</u>	Jun 03, 2013
<u>AB</u>		<u>80MG</u>	<u>A202104 006</u>	Jun 03, 2013
<u>AB</u>		<u>100MG</u>	<u>A202104 007</u>	Jun 03, 2013
	KADIAN			
	+ ALLERGAN	130MG	N020616 011	Jul 09, 2012
	+	150MG	N020616 012	Jul 09, 2012
	+!	200MG	N020616 007	Feb 27, 2007
	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

INJECTABLE; INJECTION

DURAMORPH PF

<u>AP</u>	+!	HIKMA	<u>0.5MG/ML</u>	<u>N018565 001</u>	Sep 18, 1984
<u>AP</u>	+!		<u>1MG/ML</u>	<u>N018565 002</u>	Sep 18, 1984

INFUMORPH

<u>AP</u>	+!	HIKMA	<u>10MG/ML</u>	<u>N018565 003</u>	Jul 19, 1991
<u>AP</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

MORPHINE SULFATE

<u>AP</u>		HIKMA	<u>4MG/ML</u>	<u>A205758 001</u>	May 21, 2015
<u>AP</u>			<u>8MG/ML</u>	<u>A205758 002</u>	May 21, 2015
<u>AP</u>			<u>10MG/ML</u>	<u>A205758 003</u>	May 21, 2015
<u>AP</u>		HOSPIRA	<u>0.5MG/ML</u>	<u>A071849 001</u>	May 11, 1988
<u>AP</u>			<u>0.5MG/ML</u>	<u>A073509 001</u>	Sep 30, 1992
<u>AP</u>			<u>1MG/ML</u>	<u>A071850 001</u>	May 11, 1988
<u>AP</u>			<u>1MG/ML</u>	<u>A073510 001</u>	Sep 30, 1992
<u>AP</u>	+!	HOSPIRA INC	<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
<u>AP</u>	+!	ICU MEDICAL INC	<u>1MG/ML</u>	<u>N019916 001</u>	Oct 30, 1992
	+!	HOSPIRA INC	2MG/ML	N202515 001	Nov 14, 2011
	+!	MERIDIAN MEDCL TECHN	15MG/ML	N019999 001	Jul 12, 1990

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
	+!		5MG/ML (5MG/ML)	N204223 003	Oct 30, 2013
	+!		8MG/ML (8MG/ML)	N204223 004	Oct 30, 2013
	+!		10MG/ML (10MG/ML)	N204223 005	Oct 30, 2013

SOLUTION; ORAL

MORPHINE SULFATE

<u>AA</u>		ANI PHARMS INC	<u>10MG/5ML</u>	<u>A205509 001</u>	Apr 17, 2018
<u>AA</u>			<u>20MG/5ML</u>	<u>A205509 002</u>	Apr 17, 2018
<u>AA</u>			<u>100MG/5ML</u>	<u>A205509 003</u>	Apr 17, 2018

PRESCRIPTION DRUG PRODUCT LIST

MORPHINE SULFATE

SOLUTION;ORAL

MORPHINE SULFATE

<u>AA</u>	HI TECH	<u>100MG/5ML</u>	<u>A208809 001</u>	Jul 06, 2017
<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>	NOSTRUM LABS INC	<u>10MG/5ML</u>	<u>A201011 001</u>	Feb 05, 2014
<u>AA</u>		<u>20MG/5ML</u>	<u>A201011 002</u>	Feb 05, 2014
<u>AA</u>		<u>100MG/5ML</u>	<u>A201011 003</u>	Oct 06, 2016
<u>AA</u>	PADDOCK LLC	<u>100MG/5ML</u>	<u>A201574 001</u>	Aug 06, 2012
<u>AA</u>	PHARM ASSOC	<u>100MG/5ML</u>	<u>A206573 001</u>	Nov 14, 2016
<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>20MG/5ML</u>	<u>A203519 001</u>	May 18, 2016
<u>AA</u>		<u>100MG/5ML</u>	<u>A203518 002</u>	May 12, 2015

TABLET;ORAL

MORPHINE SULFATE

<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>	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>	MAYNE PHARMA INC	<u>15MG</u>	<u>A205386 001</u>	Oct 28, 2016
<u>AB</u>		<u>30MG</u>	<u>A205386 002</u>	Oct 28, 2016
<u>AB</u>		<u>60MG</u>	<u>A205386 003</u>	Oct 28, 2016
<u>AB</u>		<u>100MG</u>	<u>A205386 004</u>	Oct 28, 2016
<u>AB</u>	NESHER PHARMS	<u>15MG</u>	<u>A076733 001</u>	May 19, 2004
<u>AB</u>		<u>30MG</u>	<u>A076720 002</u>	Dec 23, 2005
<u>AB</u>		<u>60MG</u>	<u>A076720 001</u>	May 19, 2004
<u>AB</u>		<u>100MG</u>	<u>A077855 001</u>	Sep 27, 2007
<u>AB</u>		<u>200MG</u>	<u>A077855 002</u>	Sep 27, 2007
<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
<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
<u>AB</u>	VINTAGE PHARMS LLC	<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

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

PRESCRIPTION DRUG PRODUCT LIST

MORPHINE SULFATE

TABLET, EXTENDED RELEASE;ORAL

MS CONTIN

AB	+		200MG	N019516 005	Nov 08, 1993
		MORPHABOND ER			
	+	DAIICHI SANKYO INC	15MG	N206544 001	Oct 02, 2015
	+		30MG	N206544 002	Oct 02, 2015
	+		60MG	N206544 003	Oct 02, 2015
	+	!	100MG	N206544 004	Oct 02, 2015

MOXIDECTIN

TABLET;ORAL

MOXIDECTIN

	+	!	MDGH	2MG	N210867 001	Jun 13, 2018
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MOXIFLOXACIN HYDROCHLORIDE

SOLUTION;INTRAVENOUS

MOXIFLOXACIN HYDROCHLORIDE

	+	!	FRESENIUS KABI USA	EQ 400MG BASE/250ML (EQ 1.6MG BASE/ML)	N205572 001	Apr 03, 2015
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MOXIFLOXACIN HYDROCHLORIDE IN SODIUM CHLORIDE 0.8% IN PLASTIC CONTAINER

	!	!	MYLAN LABS LTD	400MG/250ML (1.6MG/ML)	A205833 001	May 05, 2017
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SOLUTION/DROPS;OPHTHALMIC

MOXIFLOXACIN HYDROCHLORIDE

AT1			AKORN	EQ 0.5% BASE	A202916 001	Nov 09, 2017
AT1			ALEMBIC PHARMS LTD	EQ 0.5% BASE	A209469 001	Feb 13, 2019
AT1			APOTEX	EQ 0.5% BASE	A090080 001	Jun 30, 2017
AT1			AUROBINDO PHARMA LTD	EQ 0.5% BASE	A206242 001	Oct 04, 2017
AT1			LUPIN LTD	EQ 0.5% BASE	A202867 001	Sep 04, 2014
AT1			WATSON LABS INC	EQ 0.5% BASE	A202525 001	Mar 06, 2015

VIGAMOX

AT1	+	!	NOVARTIS	EQ 0.5% BASE	N021598 001	Apr 15, 2003
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MOXEZA

AT2	+	!	NOVARTIS	EQ 0.5% BASE	N022428 001	Nov 19, 2010
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MOXIFLOXACIN HYDROCHLORIDE

AT2			LUPIN LTD	EQ 0.5% BASE	A204079 001	May 28, 2015
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TABLET;ORAL

AVELOX

AB	+	!	BAYER HLTHCARE	EQ 400MG BASE	N021085 001	Dec 10, 1999
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MOXIFLOXACIN HYDROCHLORIDE

AB			AUROBINDO PHARMA LTD	EQ 400MG BASE	A202632 001	Mar 04, 2014
AB			CROSSMEDIKA SA	EQ 400MG BASE	A205348 001	Jan 14, 2016
AB			DR REDDYS LABS LTD	EQ 400MG BASE	A076938 001	Mar 04, 2014
AB			MSN	EQ 400MG BASE	A208682 001	Sep 22, 2017
AB			NOVEL LABS INC	EQ 400MG BASE	A207285 001	Feb 13, 2017
AB			TEVA PHARMS USA	EQ 400MG BASE	A077437 001	Feb 18, 2014
AB			TORRENT	EQ 400MG BASE	A200160 001	Apr 03, 2014

MUPIROCIIN

OINTMENT;TOPICAL

MUPIROCIIN

AB			FOUGERA PHARMS	2%	A065192 001	Nov 30, 2005
AB			GLENMARK PHARMS	2%	A090480 001	Jun 08, 2011
AB	!		PERRIGO ISRAEL	2%	A065123 001	Nov 07, 2003
AB			TARO	2%	A065170 001	Sep 23, 2005
AB			TEVA	2%	A065085 001	Nov 07, 2003
			CENTANY			
BX			PERRIGO NEW YORK	2%	N050788 001	Dec 04, 2002

MUPIROCIIN CALCIUM

CREAM;TOPICAL

MUPIROCIIN

	!		GLENMARK PHARMS INC	EQ 2% BASE	A201587 001	Jan 24, 2013
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MYCOPHENOLATE MOFETIL

CAPSULE;ORAL

CELLCEPT

AB	+	!	ROCHE PALO	250MG	N050722 001	May 03, 1995
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MYCOPHENOLATE MOFETIL

AB			ACCORD HLTHCARE	250MG	A090253 001	May 04, 2009
AB			ALKEM LABS LTD	250MG	A200197 001	Jun 13, 2013
AB			CONCORD BIOTECH LTD	250MG	A210181 001	Jan 08, 2019
AB			HIKMA	250MG	A065410 001	Jul 29, 2008
AB			MYLAN	250MG	A065520 001	May 04, 2009

PRESCRIPTION DRUG PRODUCT LIST

MYCOPHENOLATE MOFETIL

CAPSULE; ORAL

MYCOPHENOLATE MOFETIL

<u>AB</u>	SANDOZ	<u>250MG</u>	<u>A065379 001</u>	Oct 15, 2008
<u>AB</u>	STRIDES PHARMA	<u>250MG</u>	<u>A090055 001</u>	Jun 10, 2010
<u>AB</u>	TEVA PHARMS	<u>250MG</u>	<u>A065491 001</u>	May 06, 2009
<u>AB</u>	VINTAGE PHARMS LLC	<u>250MG</u>	<u>A090111 001</u>	Dec 22, 2009
<u>AB</u>	ZHEJIANG HISUN PHARM	<u>250MG</u>	<u>A204077 001</u>	Nov 13, 2017

SUSPENSION; ORAL

CELLCEPT

<u>AB</u>	<u>+</u> ! ROCHE PALO	<u>200MG/ML</u>	<u>N050759 001</u>	Oct 01, 1998
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MYCOPHENOLATE MOFETIL

<u>AB</u>	ALKEM LABS LTD	<u>200MG/ML</u>	<u>A203005 001</u>	Nov 14, 2014
<u>AB</u>	VISTAPHARM	<u>200MG/ML</u>	<u>A210370 001</u>	Feb 12, 2019

TABLET; ORAL

CELLCEPT

<u>AB</u>	<u>+</u> ! ROCHE PALO	<u>500MG</u>	<u>N050723 001</u>	Jun 19, 1997
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MYCOPHENOLATE MOFETIL

<u>AB</u>	ACCORD HLTHCARE	<u>500MG</u>	<u>A065416 001</u>	May 04, 2009
<u>AB</u>	ALKEM LABS LTD	<u>500MG</u>	<u>A091249 001</u>	Nov 04, 2011
<u>AB</u>	HIKMA	<u>500MG</u>	<u>A065413 001</u>	Jul 29, 2008
<u>AB</u>	MYLAN	<u>500MG</u>	<u>A065521 001</u>	May 04, 2009
<u>AB</u>	SANDOZ	<u>500MG</u>	<u>A065451 001</u>	Oct 15, 2008
<u>AB</u>	STRIDES PHARMA	<u>500MG</u>	<u>A090456 001</u>	Jun 10, 2010
<u>AB</u>	ZHEJIANG HISUN PHARM	<u>500MG</u>	<u>A204076 001</u>	Nov 16, 2017

MYCOPHENOLATE MOFETIL HYDROCHLORIDE

INJECTABLE; INJECTION

CELLCEPT

<u>AP</u>	<u>+</u> ! ROCHE PALO	<u>500MG/VIAL</u>	<u>N050758 001</u>	Aug 12, 1998
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MYCOPHENOLATE MOFETIL HYDROCHLORIDE

<u>AP</u>	AKORN INC	<u>500MG/VIAL</u>	<u>A204043 001</u>	Feb 28, 2017
<u>AP</u>	MYLAN LABS LTD	<u>500MG/VIAL</u>	<u>A203859 001</u>	Mar 31, 2017
<u>AP</u>	PAR STERILE PRODUCTS	<u>500MG/VIAL</u>	<u>A203575 001</u>	Oct 28, 2016
<u>AP</u>	ZYDUS PHARMS	<u>500MG/VIAL</u>	<u>A204473 001</u>	Aug 31, 2017

MYCOPHENOLIC ACID

TABLET, DELAYED RELEASE; ORAL

MYCOPHENOLIC ACID

<u>AB</u>	ACCORD HLTHCARE	<u>180MG</u>	<u>A202555 001</u>	Aug 23, 2017
<u>AB</u>		<u>360MG</u>	<u>A202555 002</u>	Aug 23, 2017
<u>AB</u>	APOTEX INC	<u>180MG</u>	<u>A091558 001</u>	Aug 21, 2012
<u>AB</u>		<u>360MG</u>	<u>A091558 002</u>	Aug 19, 2014
<u>AB</u>	CONCORD BIOTECH LTD	<u>180MG</u>	<u>A211173 001</u>	Dec 13, 2019
<u>AB</u>		<u>360MG</u>	<u>A211173 002</u>	Dec 13, 2019
<u>AB</u>	MYLAN	<u>180MG</u>	<u>A091248 002</u>	Jan 08, 2014
<u>AB</u>		<u>360MG</u>	<u>A091248 001</u>	Jan 08, 2014
<u>AB</u>	TEVA PHARMS USA	<u>180MG</u>	<u>A202720 001</u>	Oct 30, 2014
<u>AB</u>		<u>360MG</u>	<u>A202720 002</u>	Oct 30, 2014

MYFORTIC

<u>AB</u>	<u>+</u> NOVARTIS	<u>180MG</u>	<u>N050791 001</u>	Feb 27, 2004
<u>AB</u>	<u>+</u> !	<u>360MG</u>	<u>N050791 002</u>	Feb 27, 2004

NABILONE

CAPSULE; ORAL

CESAMET

<u>+</u> !	MYLAN SPECIALITY LP	1MG	N018677 001	Dec 26, 1985
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NABUMETONE

TABLET; ORAL

NABUMETONE

<u>AB</u>	CASI PHARMS INC	<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>	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>	IMPAX LABS INC	<u>500MG</u>	<u>A075189 001</u>	May 26, 2000
<u>AB</u>	!	<u>750MG</u>	<u>A075189 002</u>	Sep 24, 2001
<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>	LUPIN LTD	<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>	MYLAN PHARMS INC	<u>500MG</u>	<u>A090516 001</u>	Jul 12, 2010

PRESCRIPTION DRUG PRODUCT LIST

NABUMETONE

TABLET; ORAL

NABUMETONE

<u>AB</u>		<u>750MG</u>	<u>A090516 002</u>	Jul 12, 2010
<u>AB</u>	NEXGEN PHARMA INC	<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>	NOSTRUM LABS INC	<u>500MG</u>	<u>A090427 001</u>	Dec 30, 2011
<u>AB</u>		<u>750MG</u>	<u>A090427 002</u>	Dec 30, 2011
<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
	NEXGEN PHARMA INC	1GM	A203166 003	Aug 30, 2019

NADOLOL

TABLET; ORAL

CORGARD

<u>AB</u>	+	US WORLDMEDS LLC	<u>20MG</u>	<u>N018063 005</u>	Oct 28, 1986
<u>AB</u>	+		<u>40MG</u>	<u>N018063 001</u>	
<u>AB</u>	+		<u>80MG</u>	<u>N018063 002</u>	

NADOLOL

<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 LTD	<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>		HERITAGE PHARMA	<u>20MG</u>	<u>A074229 001</u>	Aug 30, 1996
<u>AB</u>			<u>40MG</u>	<u>A074229 002</u>	Aug 30, 1996
<u>AB</u>			<u>80MG</u>	<u>A074255 001</u>	Jan 24, 1996
<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>		LUPIN LTD	<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>		MYLAN	<u>20MG</u>	<u>A074172 001</u>	Oct 31, 1993
<u>AB</u>			<u>40MG</u>	<u>A074172 002</u>	Oct 31, 1993
<u>AB</u>			<u>80MG</u>	<u>A074172 003</u>	Oct 31, 1993
<u>AB</u>		NOVAST LABS	<u>20MG</u>	<u>A210786 001</u>	Jun 01, 2018
<u>AB</u>			<u>40MG</u>	<u>A210786 002</u>	Jun 01, 2018
<u>AB</u>			<u>80MG</u>	<u>A210786 003</u>	Jun 01, 2018
<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>		VGYAAN	<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>		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

+! GD SEARLE LLC

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>		AUROBINDO PHARMA LTD	<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 1GM BASE/VIAL</u>	<u>A206682 001</u>	Dec 10, 2019
<u>AP</u>			<u>EQ 2GM BASE/VIAL</u>	<u>A206682 002</u>	Dec 10, 2019
<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

PRESCRIPTION DRUG PRODUCT LIST

NAFCILLIN SODIUM

INJECTABLE; INJECTION

NAFCILLIN SODIUM

<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>	! SANDOZ	<u>EQ 1GM BASE/VIAL</u>	<u>A062527 002</u>	Aug 02, 1984
<u>AP</u>	!	<u>EQ 1GM BASE/VIAL</u>	<u>A062732 001</u>	Dec 23, 1986
<u>AP</u>	!	<u>EQ 2GM BASE/VIAL</u>	<u>A062527 003</u>	Aug 02, 1984
<u>AP</u>	!	<u>EQ 2GM BASE/VIAL</u>	<u>A062732 002</u>	Dec 23, 1986
<u>AP</u>	!	<u>EQ 10GM BASE/VIAL</u>	<u>A062527 004</u>	Aug 02, 1984
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>	TARO PHARMS	<u>2%</u>	<u>A206901 001</u>	Jan 06, 2016
<u>AB</u>	TOLMAR	<u>2%</u>	<u>A206960 001</u>	Apr 10, 2017
<u>NAFTIN</u>				
<u>AB</u>	+! SEBELA IRELAND LTD	<u>2%</u>	<u>N019599 002</u>	Jan 13, 2012
NAFTIFINE HYDROCHLORIDE				
	! TARO PHARMS	1%	A205975 001	Sep 08, 2016
GEL; TOPICAL				
<u>NAFTIFINE HYDROCHLORIDE</u>				
<u>AB</u>	TOLMAR	<u>1%</u>	<u>A206165 001</u>	Mar 20, 2019
<u>NAFTIN</u>				
<u>AB</u>	+! SEBELA IRELAND LTD	<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

NALDEMEDINE TOSYLATE

TABLET; ORAL

SYMPROIC

	+! BDSI	EQ 0.2MG BASE	N208854 001	Mar 23, 2017
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NALOXEGOL OXALATE

TABLET; ORAL

MOVANTIK

	+ ASTRAZENECA PHARMS	EQ 12.5MG BASE	N204760 001	Sep 16, 2014
	+!	EQ 25MG BASE	N204760 002	Sep 16, 2014

NALOXONE HYDROCHLORIDE

INJECTABLE; INJECTION

NALOXONE

<u>AP</u>	WEST-WARD PHARMS INT	<u>0.4MG/ML</u>	<u>A070299 001</u>	Sep 24, 1986
<u>NALOXONE HYDROCHLORIDE</u>				
<u>AP</u>	AKORN	<u>0.4MG/ML</u>	<u>A208871 001</u>	Feb 28, 2017
<u>AP</u>		<u>0.4MG/ML</u>	<u>A208872 001</u>	Mar 14, 2017
<u>AP</u>	AUROBINDO PHARMA LTD	<u>0.4MG/ML</u>	<u>A212455 001</u>	Oct 15, 2019
<u>AP</u>		<u>0.4MG/ML</u>	<u>A212456 001</u>	Nov 04, 2019
<u>AP</u>	DR REDDYS	<u>1MG/ML</u>	<u>A213209 001</u>	Mar 16, 2020
<u>AP</u>	! HOSPIRA	<u>0.4MG/ML</u>	<u>A070172 001</u>	Sep 24, 1986
<u>AP</u>	!	<u>0.4MG/ML</u>	<u>A070254 001</u>	Jan 07, 1987
<u>AP</u>	!	<u>0.4MG/ML</u>	<u>A070256 001</u>	Jan 07, 1987
<u>AP</u>	!	<u>0.4MG/ML</u>	<u>A070257 001</u>	Jan 07, 1987
<u>AP</u>	INTL MEDICATION	<u>0.4MG/ML</u>	<u>A070639 001</u>	Sep 24, 1986
<u>AP</u>	!	<u>1MG/ML</u>	<u>A072076 001</u>	Mar 24, 1988
<u>AP</u>	MYLAN INSTITUTIONAL	<u>0.4MG/ML</u>	<u>A204997 001</u>	Mar 06, 2014
<u>AP</u>		<u>0.4MG/ML</u>	<u>A205014 001</u>	Jun 29, 2016
<u>AP</u>	PAR STERILE PRODUCTS	<u>0.4MG/ML</u>	<u>A211286 001</u>	Jan 17, 2020
<u>AP</u>	SOMERSET THERAPS LLC	<u>0.4MG/ML</u>	<u>A207633 001</u>	Aug 08, 2017
<u>AP</u>		<u>0.4MG/ML</u>	<u>A207634 001</u>	Jul 26, 2017

PRESCRIPTION DRUG PRODUCT LIST

NALOXONE HYDROCHLORIDE

SOLUTION;INTRAMUSCULAR, SUBCUTANEOUS

EVZIO (AUTOINJECTOR)

+! KALEO INC 2MG/0.4ML (2MG/0.4ML)

N209862 001 Oct 19, 2016

SPRAY, METERED;NASAL

NARCAN

+! ADAPT 4MG/SPRAY

N208411 001 Nov 18, 2015

NALOXONE HYDROCHLORIDE; PENTAZOCINE HYDROCHLORIDE

TABLET;ORAL

NALOXONE HYDROCHLORIDE AND PENTAZOCINE HYDROCHLORIDE

AB	LUPIN	EQ 0.5MG BASE;EQ 50MG BASE	A075735 001	Jul 11, 2001
AB	SUN PHARM INDS LTD	EQ 0.5MG BASE;EQ 50MG BASE	A075523 001	Mar 17, 2000
AB	! WATSON LABS	EQ 0.5MG BASE;EQ 50MG BASE	A074736 001	Jan 21, 1997

NALTREXONE

FOR SUSPENSION, EXTENDED RELEASE;INTRAMUSCULAR

VIVITROL

+! ALKERMES 380MG/VIAL

N021897 001 Apr 13, 2006

NALTREXONE HYDROCHLORIDE

TABLET;ORAL

NALTREXONE HYDROCHLORIDE

AB	ACCORD HLTHCARE	50MG	A091205 001	Aug 17, 2011
AB	APOTEX	50MG	A207905 001	Jul 21, 2017
AB	BARR	50MG	A074918 001	May 08, 1998
AB	ELITE LABS	50MG	A075274 001	May 26, 1999
AB	! SPECGX LLC	50MG	A076264 002	Mar 22, 2002
AB	SUN PHARM	50MG	A090356 001	Feb 24, 2012
	SPECGX LLC	25MG	A076264 001	Mar 22, 2002
		100MG	A076264 003	Mar 22, 2002

NAPROXEN

SUSPENSION;ORAL

NAPROSYN

AB	+ ATNAHS PHARMA US	25MG/ML	N018965 001	Mar 23, 1987
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NAPROXEN

AB	! HIKMA	25MG/ML	A074190 001	Mar 30, 1994
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TABLET;ORAL

NAPROSYN

AB	+! ATNAHS PHARMA US	500MG	N017581 004	Apr 15, 1982
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NAPROXEN

AB	AMNEAL PHARMS NY	250MG	A075927 001	Dec 18, 2001
AB		375MG	A075927 002	Dec 18, 2001
AB		500MG	A075927 003	Dec 18, 2001
AB	AUROBINDO PHARMA LTD	250MG	A200429 001	Nov 08, 2011
AB		375MG	A200429 002	Nov 08, 2011
AB		500MG	A200429 003	Nov 08, 2011
AB	GLENMARK GENERICS	250MG	A078250 001	Mar 28, 2007
AB		375MG	A078250 002	Mar 28, 2007
AB		500MG	A078250 003	Mar 28, 2007
AB	INVAGEN PHARMS	250MG	A091305 001	Aug 24, 2011
AB		375MG	A091305 002	Aug 24, 2011
AB		500MG	A091305 003	Aug 24, 2011
AB	MARKSANS PHARMA	250MG	A091416 001	Feb 14, 2011
AB		375MG	A091416 002	Feb 14, 2011
AB		500MG	A091416 003	Feb 14, 2011
AB	PERRIGO PHARMS CO	250MG	A077339 001	Apr 27, 2005
AB		375MG	A077339 002	Apr 27, 2005
AB		500MG	A077339 003	Apr 27, 2005
AB	SCIEGEN PHARMS INC	250MG	A212517 001	Feb 21, 2020
AB		375MG	A212517 002	Feb 21, 2020
AB		500MG	A212517 003	Feb 21, 2020
AB	TEVA	250MG	A074201 001	Dec 21, 1993
AB		375MG	A074201 002	Dec 21, 1993
AB		500MG	A074201 003	Dec 21, 1993
AB	ZYDUS PHARMS USA	250MG	A078620 001	Jun 07, 2007
AB		375MG	A078620 002	Jun 07, 2007
AB		500MG	A078620 003	Jun 07, 2007

TABLET, DELAYED RELEASE;ORAL

EC-NAPROSYN

AB	+! ATNAHS PHARMA US	375MG	N020067 002	Oct 14, 1994
AB	+!	500MG	N020067 003	Oct 14, 1994

PRESCRIPTION DRUG PRODUCT LIST

NAPROXEN

TABLET, DELAYED RELEASE;ORAL

NAPROXEN

<u>AB</u>	PLIVA	<u>375MG</u>	<u>A075337 001</u>	May 26, 1999
<u>AB</u>		<u>500MG</u>	<u>A075337 002</u>	May 26, 1999
<u>AB</u>	SUNRISE PHARM 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

NAPROXEN SODIUM

TABLET;ORAL

ANAPROX DS

<u>AB</u>	+!	ATNAHS PHARMA US	<u>EQ 500MG BASE</u>	<u>N018164 003</u>	Sep 30, 1987
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NAPROXEN SODIUM

<u>AB</u>		AMNEAL PHARMS NY	<u>EQ 250MG BASE</u>	<u>A078432 001</u>	Apr 25, 2007
<u>AB</u>			<u>EQ 500MG BASE</u>	<u>A078432 002</u>	Apr 25, 2007
<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
<u>AB</u>		TEVA	<u>EQ 250MG BASE</u>	<u>A074198 001</u>	Dec 21, 1993
<u>AB</u>			<u>EQ 500MG BASE</u>	<u>A074198 002</u>	Dec 21, 1993

TABLET, EXTENDED RELEASE;ORAL

NAPRELAN

<u>AB</u>	+	ALVOGEN	<u>EQ 375MG BASE</u>	<u>N020353 001</u>	Jan 05, 1996
<u>AB</u>	+		<u>EQ 500MG BASE</u>	<u>N020353 002</u>	Jan 05, 1996
<u>AB</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>		MYLAN	<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>	+!	CURRAX	<u>500MG;EQ 85MG BASE</u>	<u>N021926 001</u>	Apr 15, 2008
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NARATRIPTAN HYDROCHLORIDE

TABLET;ORAL

AMERGE

<u>AB</u>	+	GLAXOSMITHKLINE LLC	<u>EQ 1MG BASE</u>	<u>N020763 002</u>	Feb 10, 1998
<u>AB</u>	+!		<u>EQ 2.5MG BASE</u>	<u>N020763 001</u>	Feb 10, 1998

NARATRIPTAN

<u>AB</u>		HERITAGE PHARMS INC	<u>EQ 1MG BASE</u>	<u>A200502 001</u>	Feb 28, 2011
<u>AB</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>		ORCHID HLTHCARE	<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>		PADDOCK LLC	<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
<u>AB</u>		SUN PHARM INDS LTD	<u>EQ 2.5MG BASE</u>	<u>A091552 001</u>	Feb 14, 2011

NATAMYCIN

SUSPENSION;OPHTHALMIC

NATACYN

	+!	NOVARTIS	5%	N050514 001	
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PRESCRIPTION DRUG PRODUCT LIST

NATEGLINIDE

TABLET; ORAL

NATEGLINIDE

<u>AB</u>	ALVOGEN	<u>60MG</u>	<u>A205055 001</u>	Dec 11, 2015
<u>AB</u>		<u>120MG</u>	<u>A205055 002</u>	Dec 11, 2015
<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>	PAR PHARM	<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>	WILSHIRE PHARMS INC	<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>	ZYDUS PHARMS	<u>60MG</u>	<u>A205248 001</u>	Jul 06, 2016
<u>AB</u>	!	<u>120MG</u>	<u>A205248 002</u>	Jul 06, 2016

NEBIVOLOL HYDROCHLORIDE

TABLET; ORAL

BYSTOLIC

<u>AB</u>	+	ALLERGAN	<u>EQ 2.5MG BASE</u>	<u>N021742 002</u>	Dec 17, 2007
<u>AB</u>	+		<u>EQ 5MG BASE</u>	<u>N021742 003</u>	Dec 17, 2007
<u>AB</u>	+		<u>EQ 10MG BASE</u>	<u>N021742 004</u>	Dec 17, 2007
<u>AB</u>	+	!	<u>EQ 20MG BASE</u>	<u>N021742 005</u>	Oct 08, 2008

NEDOCROMIL SODIUM

SOLUTION/DROPS; OPHTHALMIC

ALOCRIIL

<u>AT</u>	+	!	ALLERGAN	<u>2%</u>	<u>N021009 001</u>	Dec 08, 1999
<u>AT</u>			<u>NEDOCROMIL SODIUM</u>			
<u>AT</u>			AKORN	<u>2%</u>	<u>A090638 001</u>	Aug 22, 2012

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

	+	!	NOVARTIS	250MG/50ML (5MG/ML)	N021877 001	Oct 28, 2005
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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

<u>AA</u>			BRECKENRIDGE	<u>500MG</u>	<u>A065468 001</u>	Mar 29, 2010
<u>AA</u>	!		TEVA	<u>500MG</u>	<u>A060304 001</u>	
<u>AA</u>			XGEN PHARMS	<u>500MG</u>	<u>A065220 001</u>	Jul 28, 2006

NEOMYCIN SULFATE; POLYMYXIN B SULFATE

SOLUTION; IRRIGATION

NEOMYCIN AND POLYMYXIN B SULFATE

<u>AT</u>			XGEN PHARMS	<u>EQ 40MG BASE/ML; 200,000 UNITS/ML</u>	<u>A065106 001</u>	Jan 31, 2006
<u>AT</u>				<u>EQ 40MG BASE/ML; 200,000 UNITS/ML</u>	<u>A065108 001</u>	Jan 31, 2006

NEOSPORIN G.U. IRRIGANT

<u>AT</u>	!		MONARCH PHARMS	<u>EQ 40MG BASE/ML; 200,000 UNITS/ML</u>	<u>A060707 001</u>	
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NEOSTIGMINE METHYLSULFATE

SOLUTION; INTRAVENOUS

BLOXIVERZ

<u>AP</u>	+	!	AVADEL LEGACY	<u>5MG/10ML (0.5MG/ML)</u>	<u>N204078 001</u>	May 31, 2013
<u>AP</u>	+	!		<u>10MG/10ML (1MG/ML)</u>	<u>N204078 002</u>	May 31, 2013

NEOSTIGMINE METHYLSULFATE

<u>AP</u>			AM REGENT	<u>5MG/10ML (0.5MG/ML)</u>	<u>A209182 001</u>	May 04, 2018
<u>AP</u>				<u>10MG/10ML (1MG/ML)</u>	<u>A209182 002</u>	May 04, 2018

PRESCRIPTION DRUG PRODUCT LIST

NEOSTIGMINE METHYLSULFATE

SOLUTION; INTRAVENOUS

NEOSTIGMINE METHYLSULFATE

<u>AP</u>	AMNEAL	<u>5MG/10ML (0.5MG/ML)</u>	<u>A210051 001</u>	Jun 15, 2018
<u>AP</u>		<u>10MG/10ML (1MG/ML)</u>	<u>A210051 002</u>	Jun 15, 2018
<u>AP</u>	AMPHASTAR PHARMS INC	<u>5MG/10ML (0.5MG/ML)</u>	<u>A209933 001</u>	Sep 25, 2017
<u>AP</u>		<u>10MG/10ML (1MG/ML)</u>	<u>A209933 002</u>	Sep 25, 2017
<u>AP</u>	AMRING PHARMS	<u>5MG/10ML (0.5MG/ML)</u>	<u>A210989 001</u>	Aug 22, 2018
<u>AP</u>		<u>10MG/10ML (1MG/ML)</u>	<u>A210989 002</u>	Aug 22, 2018
<u>AP</u>	BE PHARMS	<u>5MG/10ML (0.5MG/ML)</u>	<u>A212512 001</u>	May 13, 2019
<u>AP</u>		<u>10MG/10ML (1MG/ML)</u>	<u>A212512 002</u>	May 13, 2019
<u>AP</u>	DR REDDYS	<u>5MG/10ML (0.5MG/ML)</u>	<u>A209135 001</u>	Jul 10, 2018
<u>AP</u>		<u>10MG/10ML (1MG/ML)</u>	<u>A209135 002</u>	Jul 10, 2018
<u>AP</u>	EUROHLTH INTL SARL	<u>5MG/10ML (0.5MG/ML)</u>	<u>A207042 001</u>	Dec 28, 2015
<u>AP</u>		<u>10MG/10ML (1MG/ML)</u>	<u>A207042 002</u>	Dec 28, 2015
<u>AP</u>	GLAND PHARMA LTD	<u>5MG/10ML (0.5MG/ML)</u>	<u>A212968 001</u>	Oct 16, 2019
<u>AP</u>		<u>10MG/10ML (1MG/ML)</u>	<u>A212968 002</u>	Oct 16, 2019
<u>AP</u>	PAR STERILE PRODUCTS	<u>5MG/10ML (0.5MG/ML)</u>	<u>A208405 001</u>	Apr 26, 2017
<u>AP</u>		<u>10MG/10ML (1MG/ML)</u>	<u>A208405 002</u>	Apr 26, 2017
	FRESENIUS KABI USA	3MG/3ML (1MG/ML)	N203629 003	Sep 18, 2018
		5MG/10ML (0.5MG/ML)	N203629 001	Jan 08, 2015
		10MG/10ML (1MG/ML)	N203629 002	Jan 08, 2015

NEPAFENAC

SUSPENSION/DROPS; OPHTHALMIC

ILEVRO

+	NOVARTIS	0.3%	N203491 001	Oct 16, 2012
	NEVANAC			
+	NOVARTIS	0.1%	N021862 001	Aug 19, 2005

NERATINIB MALEATE

TABLET; ORAL

NERLYNX

+	PUMA BIOTECH	EQ 40MG BASE	N208051 001	Jul 17, 2017
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NETARSUDIL MESYLATE

SOLUTION/DROPS; OPHTHALMIC

RHOPRESSA

+	AERIE PHARMS INC	EQ 0.02% BASE	N208254 001	Dec 18, 2017
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NETUPITANT; PALONOSETRON HYDROCHLORIDE

CAPSULE; ORAL

AKYNZEO

+	HELSINN HLTHCARE	300MG; EQ 0.5MG BASE	N205718 001	Oct 10, 2014
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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

VIRAMUNE

<u>AA</u>	+	BOEHRINGER INGELHEIM	<u>50MG/5ML</u>	<u>N020933 001</u>	Sep 11, 1998
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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 LABS	<u>200MG</u>	<u>A078864 001</u>	May 22, 2012
<u>AB</u>	MYLAN PHARMS INC	<u>200MG</u>	<u>A202523 001</u>	May 22, 2012
<u>AB</u>	PRINSTON INC	<u>200MG</u>	<u>A078644 001</u>	May 22, 2012
<u>AB</u>	STRIDES PHARMA	<u>200MG</u>	<u>A078195 001</u>	May 22, 2012

VIRAMUNE

<u>AB</u>	+	BOEHRINGER INGELHEIM	<u>200MG</u>	<u>N020636 001</u>	Jun 21, 1996
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TABLET, EXTENDED RELEASE; ORAL

NEVIRAPINE

<u>AB</u>	ALVOGEN	<u>100MG</u>	<u>A204621 002</u>	Nov 09, 2015
<u>AB</u>		<u>400MG</u>	<u>A204621 001</u>	Jul 10, 2015
<u>AB</u>	AUROBINDO PHARMA LTD	<u>100MG</u>	<u>A208616 001</u>	Nov 23, 2016
<u>AB</u>		<u>400MG</u>	<u>A207698 001</u>	Feb 28, 2017

PRESCRIPTION DRUG PRODUCT LISTNEVIRAPINE

TABLET, EXTENDED RELEASE;ORAL

NEVIRAPINE

AB	MACLEODS PHARMS LTD	400MG	A206879 001	Oct 06, 2017
AB	MYLAN	400MG	A205651 001	Oct 27, 2014
AB	SANDOZ INC	400MG	A203411 001	Apr 03, 2014

VIRAMUNE XR

AB	+ BOEHRINGER INGELHEIM	100MG	N201152 002	Nov 08, 2012
AB	+!	400MG	N201152 001	Mar 25, 2011

NIACIN

TABLET;ORAL

NIACIN

AA	WOCKHARDT	500MG	A081134 001	Apr 28, 1992
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NIACOR

AA	! AVONDALE PHARMS	500MG	A040378 001	May 03, 2000
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TABLET, EXTENDED RELEASE;ORAL

NIACIN

AB	AMNEAL PHARMS	500MG	A203578 001	Jul 24, 2015
AB		750MG	A204178 001	Dec 11, 2015
AB		1GM	A203578 002	Jul 24, 2015
AB	AUROBINDO PHARMA LTD	500MG	A209236 001	Feb 01, 2018
AB		750MG	A209236 002	Feb 01, 2018
AB		1GM	A209236 003	Feb 01, 2018
AB	BARR	500MG	A076378 001	Apr 26, 2005
AB		750MG	A076378 002	Apr 26, 2005
AB		1GM	A076250 001	Apr 14, 2005
AB	JUBILANT GENERICS	500MG	A209156 001	May 14, 2018
AB		750MG	A209156 002	May 14, 2018
AB		1GM	A209156 003	May 14, 2018
AB	LANNETT CO INC	500MG	A203899 001	Jun 16, 2017
AB		1GM	A203899 002	Jun 16, 2017
AB	LUPIN LTD	500MG	A090860 001	Mar 20, 2014
AB		750MG	A090892 001	Mar 20, 2014
AB		1GM	A090446 001	Mar 20, 2014
AB	SUN PHARM	500MG	A200484 001	Apr 23, 2014
AB		750MG	A201273 001	Apr 23, 2014
AB		1GM	A200484 002	Apr 23, 2014

NIASPAN

AB	+ ABBVIE	500MG	N020381 002	Jul 28, 1997
AB	+!	750MG	N020381 003	Jul 28, 1997
AB	+!	1GM	N020381 004	Jul 28, 1997

NICARDIPINE HYDROCHLORIDE

CAPSULE;ORAL

NICARDIPINE HYDROCHLORIDE

AB	EPIC PHARMA	20MG	A074928 001	Mar 19, 1998
AB		30MG	A074928 002	Mar 19, 1998
AB	MYLAN	20MG	A074642 001	Jul 18, 1996
AB	!	30MG	A074642 002	Jul 18, 1996

INJECTABLE;INJECTION

NICARDIPINE HYDROCHLORIDE

EXELA PHARMA SCIENCE	25MG/10ML (2.5MG/ML)	N022276 001	Jul 24, 2008
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INJECTABLE;INTRAVENOUS

CARDENE IN 0.83% SODIUM CHLORIDE IN PLASTIC CONTAINER

+! CHIESI USA INC	40MG/200ML (0.2MG/ML)	N019734 004	Nov 07, 2008
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CARDENE IN 0.86% SODIUM CHLORIDE IN PLASTIC CONTAINER

+! CHIESI USA INC	20MG/200ML (0.1MG/ML)	N019734 003	Jul 31, 2008
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CARDENE IN 4.8% DEXTROSE IN PLASTIC CONTAINER

+! CHIESI USA INC	20MG/200ML (0.1MG/ML)	N019734 002	Jul 31, 2008
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NICOTINE

INHALANT;ORAL

NICOTROL

+! PHARMACIA AND UPJOHN	4MG/CARTRIDGE	N020714 001	May 02, 1997
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SPRAY, METERED;NASAL

NICOTROL

+! PFIZER INC	0.5MG/SPRAY	N020385 001	Mar 22, 1996
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PRESCRIPTION DRUG PRODUCT LIST

3-318 (of 453)

NIFEDIPINE

CAPSULE;ORAL

NIFEDIPINE

<u>AB</u>	ACTAVIS ELIZABETH	<u>10MG</u>	<u>A072579 001</u>	Jan 08, 1991
<u>AB</u>		<u>20MG</u>	<u>A072556 001</u>	Sep 20, 1990
<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>	INTERGEL PHARM	<u>10MG</u>	<u>A072781 001</u>	Jul 30, 1993
<u>AB</u>	LEADING PHARMA LLC	<u>10MG</u>	<u>A073250 001</u>	Oct 08, 1991
<u>AB</u>		<u>20MG</u>	<u>A074045 001</u>	Apr 30, 1992

PROCARDIA

<u>AB</u>	+! PFIZER	<u>10MG</u>	<u>N018482 001</u>	
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TABLET, EXTENDED RELEASE;ORAL

ADALAT CC

<u>AB1</u>	+ ALVOGEN	<u>30MG</u>	<u>N020198 001</u>	Apr 21, 1993
<u>AB1</u>	+!	<u>60MG</u>	<u>N020198 002</u>	Apr 21, 1993
<u>AB1</u>	+!	<u>90MG</u>	<u>N020198 003</u>	Apr 21, 1993

NIFEDIPINE

<u>AB1</u>	MYLAN	<u>30MG</u>	<u>A201071 001</u>	Dec 03, 2010
<u>AB1</u>		<u>60MG</u>	<u>A201071 002</u>	Dec 03, 2010
<u>AB1</u>		<u>90MG</u>	<u>A201071 003</u>	Dec 03, 2010
<u>AB1</u>	NOVAST LABS	<u>30MG</u>	<u>A202987 001</u>	Aug 25, 2016
<u>AB1</u>		<u>60MG</u>	<u>A202987 002</u>	Aug 25, 2016
<u>AB1</u>		<u>90MG</u>	<u>A202987 003</u>	Aug 25, 2016
<u>AB1</u>	PAR PHARM	<u>30MG</u>	<u>A077899 001</u>	Dec 13, 2006
<u>AB1</u>		<u>60MG</u>	<u>A077899 002</u>	Dec 13, 2006
<u>AB1</u>		<u>90MG</u>	<u>A077899 003</u>	May 25, 2012
<u>AB1</u>	VALEANT PHARMS NORTH	<u>30MG</u>	<u>A075269 001</u>	Dec 04, 2000
<u>AB1</u>		<u>60MG</u>	<u>A075269 002</u>	Dec 04, 2000
<u>AB1</u>		<u>90MG</u>	<u>A076070 001</u>	Aug 16, 2002
<u>AB1</u>	ZYDUS PHARMS	<u>30MG</u>	<u>A210184 001</u>	Jun 29, 2018
<u>AB1</u>		<u>60MG</u>	<u>A210184 002</u>	Jun 29, 2018
<u>AB1</u>		<u>90MG</u>	<u>A210184 003</u>	Jun 29, 2018
<u>AB2</u>	NOVAST LABS	<u>30MG</u>	<u>A210614 001</u>	Mar 12, 2019
<u>AB2</u>		<u>60MG</u>	<u>A210614 002</u>	Mar 12, 2019
<u>AB2</u>		<u>90MG</u>	<u>A210614 003</u>	Mar 12, 2019
<u>AB2</u>	OSMOTICA PHARM US	<u>30MG</u>	<u>A077127 001</u>	Nov 21, 2005
<u>AB2</u>		<u>60MG</u>	<u>A077127 002</u>	Nov 21, 2005
<u>AB2</u>		<u>90MG</u>	<u>A077127 003</u>	Oct 03, 2007
<u>AB2</u>	SPIL	<u>30MG</u>	<u>A210838 001</u>	Apr 16, 2019
<u>AB2</u>		<u>60MG</u>	<u>A210838 002</u>	Apr 16, 2019
<u>AB2</u>		<u>90MG</u>	<u>A210838 003</u>	Apr 16, 2019
<u>AB2</u>	TWI PHARMS	<u>30MG</u>	<u>A203126 001</u>	Apr 03, 2014
<u>AB2</u>		<u>60MG</u>	<u>A203126 002</u>	Apr 03, 2014
<u>AB2</u>		<u>90MG</u>	<u>A203126 003</u>	Apr 03, 2014
<u>AB2</u>	VALEANT PHARMS NORTH	<u>30MG</u>	<u>A075289 002</u>	Feb 06, 2001
<u>AB2</u>		<u>60MG</u>	<u>A075289 001</u>	Sep 27, 2000
<u>AB2</u>	ZYDUS PHARMS	<u>30MG</u>	<u>A210012 001</u>	Dec 19, 2017
<u>AB2</u>		<u>60MG</u>	<u>A210012 002</u>	Dec 19, 2017
<u>AB2</u>		<u>90MG</u>	<u>A210012 003</u>	Dec 19, 2017

PROCARDIA XL

<u>AB2</u>	+ PFIZER	<u>30MG</u>	<u>N019684 001</u>	Sep 06, 1989
<u>AB2</u>	+	<u>60MG</u>	<u>N019684 002</u>	Sep 06, 1989
<u>AB2</u>	+!	<u>90MG</u>	<u>N019684 003</u>	Sep 06, 1989

NILOTINIB HYDROCHLORIDE

CAPSULE;ORAL

TASIGNA

+	NOVARTIS	EQ 50MG BASE	N022068 003	Mar 22, 2018
+		EQ 150MG BASE	N022068 002	Jun 17, 2010
+	!	EQ 200MG BASE	N022068 001	Oct 29, 2007

NILUTAMIDE

TABLET;ORAL

NILANDRON

<u>AB</u>	+! CONCORDIA	<u>150MG</u>	<u>N020169 002</u>	Apr 30, 1999
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NILUTAMIDE

<u>AB</u>	ANI PHARMS INC	<u>150MG</u>	<u>A207631 001</u>	Jul 15, 2016
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PRESCRIPTION DRUG PRODUCT LIST

NIMODIPINE

CAPSULE; ORAL

NIMODIPINE

AB	!	BIONPHARMA INC	30MG	A076740	001	Jan 17, 2008
AB		HERITAGE PHARMS INC	30MG	A077811	001	May 02, 2007
AB		SOFGEN PHARMS	30MG	A201832	001	Jul 24, 2015
AB		THEPHARMANETWORK LLC	30MG	A090103	001	Apr 07, 2014

SOLUTION; ORAL

NYMALIZE

+	!	ARBOR PHARMS LLC	60MG/20ML	N203340	001	May 10, 2013
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NINTEDANIB ESYLATE

CAPSULE; ORAL

OFEV

+		BOEHRINGER INGELHEIM	EQ 100MG BASE	N205832	001	Oct 15, 2014
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+	!		EQ 150MG BASE	N205832	002	Oct 15, 2014
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NIRAPARIB TOSYLATE

CAPSULE; ORAL

ZEJULA

+	!	GLAXOSMITHKLINE	EQ 100MG BASE	N208447	001	Mar 27, 2017
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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 PHARMA BV	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

FOR SUSPENSION; ORAL

ALINIA

+	!	ROMARK	100MG/5ML	N021498	001	Nov 22, 2002
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TABLET; ORAL

ALINIA

+	!	ROMARK	500MG	N021497	001	Jul 21, 2004
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NITISINONE

CAPSULE; ORAL

NITISINONE

AB		NOVITIUM PHARMA	2MG	A211041	001	Aug 26, 2019
AB			5MG	A211041	002	Aug 26, 2019
AB			10MG	A211041	003	Aug 26, 2019

ORFADIN

AB	+		2MG	N021232	001	Jan 18, 2002
AB	+	SWEDISH ORPHAN	5MG	N021232	002	Jan 18, 2002
AB	+		10MG	N021232	003	Jan 18, 2002
	+	!	20MG	N021232	004	Jun 13, 2016

SUSPENSION; ORAL

ORFADIN

+	!	SWEDISH ORPHAN	4MG/ML	N206356	001	Apr 22, 2016
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TABLET; ORAL

NITYR

+		CYCLE PHARMS LTD	2MG	N209449	001	Jul 26, 2017
+			5MG	N209449	002	Jul 26, 2017
+	!		10MG	N209449	003	Jul 26, 2017

NITRIC OXIDE

GAS; INHALATION

INOMAX

AA	+	!	MALLINCKRODT HOSP	800PPM	N020845	003	Dec 23, 1999
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NOXIVENT

AA		PRAXAIR DISTRIBUTION	800PPM	A207141	002	Oct 02, 2018
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PRESCRIPTION DRUG PRODUCT LISTNITRIC OXIDE

GAS; INHALATION

GENOSYL

+! VERO BIOTECH 800PPM N202860 001 Dec 20, 2019

NOXIVENT

PRAXAIR 100PPM A207141 001 Oct 02, 2018
DISTRIBUTIONNITROFURANTOIN

SUSPENSION; ORAL

FURADANTIN**AB** +! CASPER PHARMA LLC 25MG/5ML **N009175 001**NITROFURANTOIN**AB** ACTAVIS MID 25MG/5ML **A205180 001** May 03, 2016

ATLANTIC

AB AMNEAL PHARMS 25MG/5ML **A201679 001** May 11, 2011**AB** NOSTRUM LABS INC 25MG/5ML **A201355 001** Aug 14, 2013**AB** NOVEL LABS INC 25MG/5ML **A201693 001** Sep 08, 2014NITROFURANTOIN, MACROCRYSTALLINE

CAPSULE; ORAL

MACRODANTIN**AB** + ALVOGEN 25MG **N016620 003****AB** + 50MG **N016620 001****AB** +! 100MG **N016620 002**NITROFURANTOIN**AB** ACTAVIS LABS FL INC 25MG **A091095 001** Jun 18, 2015**AB** 50MG **A091095 002** Jun 18, 2015**AB** 100MG **A091095 003** Jun 18, 2015**AB** IMPAX LABS INC 50MG **A073671 001** Jan 28, 1993**AB** 100MG **A073652 001** Jan 28, 1993**AB** NOVEL LABS INC 50MG **A203233 001** Jul 09, 2018**AB** 100MG **A203233 002** Jul 09, 2018**AB** SUN PHARM 25MG **A201722 001** Feb 16, 2016

INDUSTRIES

AB 50MG **A201722 002** Feb 16, 2016**AB** 100MG **A201722 003** Feb 16, 2016**AB** ZYDUS PHARMS 50MG **A205005 001** Dec 12, 2017**AB** 100MG **A205005 002** Dec 12, 2017NITROFURANTOIN; NITROFURANTOIN, MACROCRYSTALLINE

CAPSULE; ORAL

MACROBID**AB** +! ALVOGEN 75MG; 25MG **N020064 001** Dec 24, 1991NITROFURANTOIN (MONOHYDRATE/MACROCRYSTALS)**AB** AMNEAL PHARMS 75MG; 25MG **A207372 001** May 15, 2017**AB** MYLAN 75MG; 25MG **A076648 001** Mar 22, 2004**AB** SANDOZ 75MG; 25MG **A077066 001** Apr 05, 2005**AB** SUNNY PHARMTECH INC 75MG; 25MG **A208516 001** May 24, 2018**AB** WATSON LABS INC 75MG; 25MG **A202250 001** Jul 08, 2015NITROGLYCERIN

AEROSOL, METERED; SUBLINGUAL

NITROMIST

+! EVUS 0.4MG/SPRAY N021780 001 Nov 02, 2006

FILM, EXTENDED RELEASE; TRANSDERMAL

NITROGLYCERIN**AB2** HERCON PHARM 0.1MG/HR **A089885 002** Oct 30, 2017**AB2** 0.2MG/HR **A089884 001** Oct 30, 1998**AB2** 0.4MG/HR **A089885 001** Oct 30, 1998**AB2** 0.6MG/HR **A089886 001** Oct 30, 1998**AB2** ! MYLAN TECHNOLOGIES 0.1MG/HR **A074559 004** Feb 06, 1998**AB2** ! 0.2MG/HR **A074559 003** Aug 30, 1996**AB2** ! 0.4MG/HR **A074559 002** Aug 30, 1996**AB2** ! 0.6MG/HR **A074559 001** Aug 30, 1996

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

PRESCRIPTION DRUG PRODUCT LIST

NITROGLYCERIN

INJECTABLE; INJECTION

NITROGLYCERIN IN DEXTROSE 5%

AP	+ !	BAXTER HLTHCARE	<u>10MG/100ML</u>	<u>N019970 001</u>	Dec 29, 1989
AP	+ !		<u>20MG/100ML</u>	<u>N019970 002</u>	Dec 29, 1989
AP	+ !		<u>40MG/100ML</u>	<u>N019970 003</u>	Dec 29, 1989

NITROGLYCERIN

! AM REGENT

5MG/ML

A072034 001 May 24, 1988

OINTMENT; INTRA-ANAL

RECTIV

+! ALLERGAN

0.4%

N021359 001 Jun 21, 2011

OINTMENT; TRANSDERMAL

NITROGLYCERIN

! FOUGERA PHARMS INC

2%

A087355 001 Jul 08, 1988

POWDER; SUBLINGUAL

GONITRO

+! POHL BOSKAMP

0.4MG/PACKET

N208424 001 Jun 08, 2016

SPRAY, METERED; SUBLINGUAL

NITROGLYCERIN

AB		PERRIGO ISRAEL	<u>0.4MG/SPRAY</u>	<u>A091496 001</u>	Sep 20, 2013
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NITROLINGUAL PUMPSPRAY

AB	+ !	POHL BOSKAMP	<u>0.4MG/SPRAY</u>	<u>N018705 002</u>	Jan 10, 1997
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TABLET; SUBLINGUAL

NITROGLYCERIN

AB		ACTAVIS LABS FL INC	<u>0.3MG</u>	<u>A203693 001</u>	Oct 16, 2017
AB			<u>0.4MG</u>	<u>A203693 002</u>	Oct 16, 2017
AB			<u>0.6MG</u>	<u>A203693 003</u>	Oct 16, 2017
AB		ALVOGEN	<u>0.3MG</u>	<u>A211604 001</u>	Apr 30, 2019
AB			<u>0.4MG</u>	<u>A211604 002</u>	Apr 30, 2019
AB			<u>0.6MG</u>	<u>A211604 003</u>	Apr 30, 2019
AB		DR REDDYS	<u>0.3MG</u>	<u>A208191 001</u>	Aug 26, 2016
AB			<u>0.4MG</u>	<u>A208191 002</u>	Aug 26, 2016
AB			<u>0.6MG</u>	<u>A208191 003</u>	Aug 26, 2016
AB		GLENMARK PHARMS SA	<u>0.3MG</u>	<u>A206391 001</u>	Sep 19, 2017
AB			<u>0.4MG</u>	<u>A206391 002</u>	Sep 19, 2017
AB			<u>0.6MG</u>	<u>A206391 003</u>	Sep 19, 2017
AB		SIGMAPHARM LABS LLC	<u>0.3MG</u>	<u>A207745 001</u>	May 07, 2018
AB			<u>0.4MG</u>	<u>A207745 002</u>	May 07, 2018
AB			<u>0.6MG</u>	<u>A207745 003</u>	May 07, 2018

NITROSTAT

AB	+	PFIZER PHARMS	<u>0.3MG</u>	<u>N021134 001</u>	May 01, 2000
AB	+		<u>0.4MG</u>	<u>N021134 002</u>	May 01, 2000
AB	+ !		<u>0.6MG</u>	<u>N021134 003</u>	May 01, 2000

NIZATIDINE

CAPSULE; ORAL

NIZATIDINE

AB		DR REDDYS LABS LTD	<u>150MG</u>	<u>A077314 001</u>	Sep 15, 2005
AB			<u>300MG</u>	<u>A077314 002</u>	Sep 15, 2005
AB		GLENMARK GENERICS	<u>150MG</u>	<u>A090618 001</u>	Jul 15, 2011
AB			<u>300MG</u>	<u>A090618 002</u>	Jul 15, 2011
AB		MYLAN PHARMS INC	<u>150MG</u>	<u>A075806 001</u>	Jul 05, 2002
AB	!		<u>300MG</u>	<u>A075806 002</u>	Jul 05, 2002
AB		SANDOZ	<u>150MG</u>	<u>A076178 001</u>	Jul 05, 2002
AB			<u>300MG</u>	<u>A076178 002</u>	Jul 05, 2002
AB		WATSON LABS	<u>150MG</u>	<u>A075616 001</u>	Jul 09, 2002
AB			<u>300MG</u>	<u>A075616 002</u>	Jul 09, 2002

SOLUTION; ORAL

NIZATIDINE

! AMNEAL PHARMS

15MG/ML

A090576 001 Nov 18, 2009

NOREPINEPHRINE BITARTRATE

INJECTABLE; INJECTION

LEVOPHED

AP	+ !	HOSPIRA	<u>EQ 1MG BASE/ML</u>	<u>N007513 001</u>	
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NOREPINEPHRINE BITARTRATE

AP		AMNEAL	<u>EQ 1MG BASE/ML</u>	<u>A210839 001</u>	Dec 17, 2018
AP		BAXTER HLTHCARE CORP	<u>EQ 1MG BASE/ML</u>	<u>A040859 001</u>	Mar 27, 2012
AP		HIKMA	<u>EQ 1MG BASE/ML</u>	<u>A203662 001</u>	Nov 07, 2018
AP		MYLAN LABS LTD	<u>EQ 1MG BASE/ML</u>	<u>A211242 001</u>	Oct 04, 2018
AP		SANDOZ INC	<u>EQ 1MG BASE/ML</u>	<u>A211359 001</u>	Oct 18, 2018
AP		TEVA PHARMS USA	<u>EQ 1MG BASE/ML</u>	<u>A040455 001</u>	Mar 03, 2003
AP		WEST-WARD PHARMS	<u>EQ 1MG BASE/ML</u>	<u>A040462 001</u>	Oct 31, 2003

PRESCRIPTION DRUG PRODUCT LIST

NOREPINEPHRINE BITARTRATE

INJECTABLE; INJECTION

NOREPINEPHRINE BITARTRATE

INT

NORETHINDRONE

TABLET; ORAL-28

CAMILA**AB1** MAYNE PHARMA **0.35MG** **A076177 001** Oct 21, 2002HEATHER**AB1** GLENMARK GENERICS **0.35MG** **A090454 001** Apr 23, 2010INCASSIA**AB1** AUROBINDO PHARMA LTD **0.35MG** **A207304 001** Sep 23, 2016NOR-OD**AB1** +! TEVA BRANDED PHARM **0.35MG** **N017060 001**NORETHINDRONE**AB1** ACCORD HLTHCARE **0.35MG** **A206807 001** Dec 13, 2016**AB1** AMNEAL PHARMS **0.35MG** **A202260 001** Aug 01, 2013**AB1** LUPIN LTD **0.35MG** **A091325 001** Sep 19, 2011**AB1** MYLAN LABS LTD **0.35MG** **A201483 001** Jun 24, 2013**AB1** NOVAST LABS **0.35MG** **A202014 001** Sep 13, 2013ERRIN**AB2** MAYNE PHARMA **0.35MG** **A076225 001** Oct 21, 2002JENCYCLA**AB2** LUPIN LTD **0.35MG** **A091323 001** Mar 28, 2013NORETHINDRONE**AB2** GLENMARK GENERICS **0.35MG** **A091209 001** Jul 22, 2010**AB2** ! MYLAN LABS LTD **0.35MG** **A200980 001** Jun 12, 2013**AB2** NOVAST LABS **0.35MG** **A200961 001** Sep 13, 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 GENERICS **5MG** **A091090 001** Jul 21, 2010**AB** MYLAN LABS LTD **5MG** **A205278 001** Nov 10, 2016**AB** NOVAST LABS **5MG** **A206490 001** Nov 05, 2018NORTRIPTYLINE HYDROCHLORIDE

CAPSULE; ORAL

NORTRIPTYLINE HYDROCHLORIDE**AB** MAYNE PHARMA **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, 1995PAMELOR**AB** + SPECGX LLC **EQ 10MG BASE** **N018013 001****AB** + **EQ 25MG BASE** **N018013 002****AB** + **EQ 50MG BASE** **N018013 004****AB** +! **EQ 75MG BASE** **N018013 003**

SOLUTION; ORAL

NORTRIPTYLINE HYDROCHLORIDE**AA** ! PHARM ASSOC **EQ 10MG BASE/5ML** **A075606 001** Aug 28, 2000**AA** TARO **EQ 10MG BASE/5ML** **A077965 001** Jun 20, 2006NUSINERSEN SODIUM

SOLUTION; INTRATHECAL

SPINRAZA

+! BIOGEN IDEC 12MG/5ML (2.4MG/ML) N209531 001 Dec 23, 2016

PRESCRIPTION DRUG PRODUCT LIST

NYSTATIN

CREAM; TOPICAL

NYSTATIN

<u>AT</u>	ACP NIMBLE	<u>100,000 UNITS/GM</u>	<u>A061966 001</u>	
<u>AT</u>	ACTAVIS MID ATLANTIC	<u>100,000 UNITS/GM</u>	<u>A062949 001</u>	Jun 13, 1988
<u>AT</u>	CROWN LABS INC	<u>100,000 UNITS/GM</u>	<u>A207733 001</u>	Sep 26, 2017
<u>AT</u>	FOUGERA PHARMS	<u>100,000 UNITS/GM</u>	<u>A062129 001</u>	
<u>AT</u>	PERRIGO NEW YORK	<u>100,000 UNITS/GM</u>	<u>A062225 001</u>	
<u>AT</u>	! TARO	<u>100,000 UNITS/GM</u>	<u>A064022 001</u>	Jan 29, 1993
<u>AT</u>	TORRENT	<u>100,000 UNITS/GM</u>	<u>A212557 001</u>	Jul 24, 2019
<u>AT</u>	VINTAGE	<u>100,000 UNITS/GM</u>	<u>A065315 001</u>	May 31, 2006

OINTMENT; TOPICAL

NYSTATIN

<u>AT</u>	ACP NIMBLE	<u>100,000 UNITS/GM</u>	<u>A209114 001</u>	Oct 06, 2017
<u>AT</u>	ACTAVIS MID ATLANTIC	<u>100,000 UNITS/GM</u>	<u>A062840 001</u>	Nov 13, 1987
<u>AT</u>	CADILA	<u>100,000 UNITS/GM</u>	<u>A207767 001</u>	May 25, 2018
<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>	PERRIGO NEW YORK	<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

POWDER; TOPICAL

NYSTATIN

<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>	! MAYNE PHARMA INC	<u>100,000 UNITS/GM</u>	<u>A065203 001</u>	Jul 15, 2004
<u>AT</u>	NESHER PHARMS	<u>100,000 UNITS/GM</u>	<u>A208581 001</u>	Jun 08, 2017
<u>AT</u>	UPSHER SMITH LABS	<u>100,000 UNITS/GM</u>	<u>A065183 001</u>	May 03, 2005
<u>AT</u>	XGEN PHARMS	<u>100,000 UNITS/GM</u>	<u>A065175 001</u>	Dec 17, 2004

NYSTOP

<u>AT</u>	PADDOCK LLC	<u>100,000 UNITS/GM</u>	<u>A064118 001</u>	Aug 16, 1996
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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>	HI TECH PHARMA	<u>100,000 UNITS/ML</u>	<u>A064042 001</u>	Feb 28, 1994
<u>AA</u>	LANNETT CO INC	<u>100,000 UNITS/ML</u>	<u>A065148 001</u>	Jun 28, 2005
<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>	VISTAPHARM	<u>100,000 UNITS/ML</u>	<u>A064142 001</u>	Jun 25, 1998
<u>AA</u>		<u>100,000 UNITS/ML</u>	<u>A065422 001</u>	Mar 07, 2011
<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 PHARMS INC	<u>500,000 UNITS</u>	<u>A062474 001</u>	Dec 22, 1983
<u>AA</u>	SUN PHARM INDUSTRIES	<u>500,000 UNITS</u>	<u>A062838 001</u>	Dec 22, 1988
<u>AA</u>	! TEVA	<u>500,000 UNITS</u>	<u>A062506 001</u>	Jan 16, 1984

NYSTATIN; TRIAMCINOLONE ACETONIDE

CREAM; TOPICAL

MYKACET

<u>AT</u>	ACP NIMBLE	<u>100,000 UNITS/GM; 0.1%</u>	<u>A062367 001</u>	May 28, 1985
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NYSTATIN AND TRIAMCINOLONE ACETONIDE

<u>AT</u>	AMNEAL PHARMS LLC	<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>	PERRIGO UK FINCO	<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

MYKACET

<u>AT</u>	ACP NIMBLE	<u>100,000 UNITS/GM; 0.1%</u>	<u>A062733 001</u>	Mar 06, 1987
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NYSTATIN AND TRIAMCINOLONE ACETONIDE

<u>AT</u>	AKORN	<u>100,000 UNITS/GM; 0.1%</u>	<u>A207217 001</u>	Aug 04, 2017
<u>AT</u>	DR REDDYS	<u>100,000 UNITS/GM; 0.1%</u>	<u>A207741 001</u>	Jan 31, 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>	PERRIGO UK FINCO	<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

PRESCRIPTION DRUG PRODUCT LIST

NYSTATIN; TRIAMCINOLONE ACETONIDE

OINTMENT; TOPICAL

NYSTATIN AND TRIAMCINOLONE ACETONIDE

<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>	TELIGENT PHARMA INC	<u>100,000 UNITS/GM;0.1%</u>	<u>A208287 001</u>	Dec 30, 2016
<u>AT</u>	VITRUVIAS THERAP	<u>100000 UNITS/GM;0.1%</u>	<u>A207316 001</u>	Nov 18, 2019

OBETICHOIC ACID

TABLET; ORAL

OCALIVA

+	INTERCEPT PHARMS	5MG	N207999 001	May 27, 2016
	INC			
+	!	10MG	N207999 002	May 27, 2016

OCTREOTIDE ACETATE

INJECTABLE; INJECTION

OCTREOTIDE ACETATE

<u>AP</u>	FRESENIUS KABI USA	<u>EQ 0.2MG BASE/ML</u>	<u>A077450 001</u>	Feb 10, 2006
<u>AP</u>		<u>EQ 1MG BASE/ML</u>	<u>A077450 002</u>	Feb 10, 2006
<u>AP</u>	SAGENT PHARMS INC	<u>EQ 0.2MG BASE/ML</u>	<u>A091041 001</u>	Nov 12, 2013
<u>AP</u>		<u>EQ 1MG BASE/ML</u>	<u>A091041 002</u>	Nov 12, 2013
<u>AP</u>	TEVA PHARMS USA	<u>EQ 0.05MG BASE/ML</u>	<u>A075957 001</u>	Oct 03, 2005
<u>AP</u>		<u>EQ 0.1MG BASE/ML</u>	<u>A075957 002</u>	Oct 03, 2005
<u>AP</u>		<u>EQ 0.2MG BASE/ML</u>	<u>A075959 001</u>	Nov 21, 2005
<u>AP</u>		<u>EQ 0.5MG BASE/ML</u>	<u>A075957 003</u>	Oct 03, 2005
<u>AP</u>		<u>EQ 1MG BASE/ML</u>	<u>A075959 002</u>	Nov 21, 2005
<u>AP</u>	! WEST-WARD PHARMS	<u>EQ 0.2MG BASE/ML</u>	<u>A076330 001</u>	Apr 08, 2005
	INT			
<u>AP</u>	!	<u>EQ 1MG BASE/ML</u>	<u>A076330 002</u>	Apr 08, 2005

OCTREOTIDE ACETATE (PRESERVATIVE FREE)

<u>AP</u>	FRESENIUS KABI USA	<u>EQ 0.05MG BASE/ML</u>	<u>A077457 001</u>	Feb 10, 2006
<u>AP</u>		<u>EQ 0.1MG BASE/ML</u>	<u>A077457 002</u>	Feb 10, 2006
<u>AP</u>		<u>EQ 0.5MG BASE/ML</u>	<u>A077457 003</u>	Feb 10, 2006
<u>AP</u>	MYLAN INSTITUTIONAL	<u>EQ 0.05MG BASE/ML</u>	<u>A079198 001</u>	Feb 10, 2011
<u>AP</u>		<u>EQ 0.1MG BASE/ML</u>	<u>A079198 002</u>	Feb 10, 2011
<u>AP</u>		<u>EQ 0.5MG BASE/ML</u>	<u>A079198 003</u>	Feb 10, 2011
<u>AP</u>	SAGENT PHARMS INC	<u>EQ 0.05MG BASE/ML</u>	<u>A090834 001</u>	Nov 12, 2013
<u>AP</u>		<u>EQ 0.1MG BASE/ML</u>	<u>A090834 002</u>	Nov 12, 2013
<u>AP</u>		<u>EQ 0.5MG BASE/ML</u>	<u>A090834 003</u>	Nov 12, 2013
<u>AP</u>	! WEST-WARD PHARMS	<u>EQ 0.05MG BASE/ML</u>	<u>A076313 001</u>	Mar 28, 2005
	INT			
<u>AP</u>	!	<u>EQ 0.1MG BASE/ML</u>	<u>A076313 003</u>	Mar 28, 2005
<u>AP</u>	!	<u>EQ 0.5MG BASE/ML</u>	<u>A076313 002</u>	Mar 28, 2005

SANDOSTATIN

<u>AP</u>	! NOVARTIS	<u>EQ 0.05MG BASE/ML</u>	<u>N019667 001</u>	Oct 21, 1988
<u>AP</u>	!	<u>EQ 0.1MG BASE/ML</u>	<u>N019667 002</u>	Oct 21, 1988
<u>AP</u>	!	<u>EQ 0.2MG BASE/ML</u>	<u>N019667 004</u>	Jun 12, 1991
<u>AP</u>	!	<u>EQ 0.5MG BASE/ML</u>	<u>N019667 003</u>	Oct 21, 1988
<u>AP</u>	!	<u>EQ 1MG BASE/ML</u>	<u>N019667 005</u>	Jun 12, 1991

SANDOSTATIN LAR

+	NOVARTIS	EQ 10MG BASE/VIAL	N021008 001	Nov 25, 1998
+		EQ 20MG BASE/VIAL	N021008 002	Nov 25, 1998
+	!	EQ 30MG BASE/VIAL	N021008 003	Nov 25, 1998

SOLUTION; SUBCUTANEOUS

BYNFEZIA PEN

+	SUN PHARM	EQ 2.5MG BASE/ML (EQ 2.5MG BASE/ML)	N213224 001	Jan 28, 2020
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OFLOXACIN

SOLUTION/DROPS; OPHTHALMIC

OCUFLOX

<u>AT</u>	! ALLERGAN	<u>0.3%</u>	<u>N019921 001</u>	Jul 30, 1993
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OFLOXACIN

<u>AT</u>	AKORN	<u>0.3%</u>	<u>A076407 001</u>	Apr 15, 2008
<u>AT</u>	ALTAIRE PHARMS INC	<u>0.3%</u>	<u>A202692 001</u>	Apr 29, 2013
<u>AT</u>	ALVOGEN	<u>0.3%</u>	<u>A076830 001</u>	Aug 31, 2004
<u>AT</u>	APOTEX INC	<u>0.3%</u>	<u>A076513 001</u>	May 14, 2004
<u>AT</u>	BAUSCH AND LOMB	<u>0.3%</u>	<u>A076622 001</u>	May 14, 2004
<u>AT</u>	FDC LTD	<u>0.3%</u>	<u>A078559 001</u>	Feb 25, 2009
<u>AT</u>	HI TECH PHARMA	<u>0.3%</u>	<u>A076615 001</u>	May 14, 2004

SOLUTION/DROPS; OTIC

OFLOXACIN

<u>AT</u>	AMNEAL	<u>0.3%</u>	<u>A211525 001</u>	Aug 30, 2019
<u>AT</u>	APOTEX INC	<u>0.3%</u>	<u>A076527 001</u>	Sep 28, 2007

PRESCRIPTION DRUG PRODUCT LIST

OFLOXACIN

SOLUTION/DROPS;OTIC

OFLOXACIN

<u>AT</u>	!	BAUSCH AND LOMB	<u>0.3%</u>	<u>A076128</u>	<u>001</u>	Mar 17, 2008
<u>AT</u>		HI TECH PHARMA	<u>0.3%</u>	<u>A076616</u>	<u>001</u>	Mar 17, 2008

TABLET;ORAL

OFLOXACIN

<u>AB</u>		CADILA PHARMS LTD	<u>200MG</u>	<u>A091656</u>	<u>001</u>	Sep 18, 2014
<u>AB</u>			<u>300MG</u>	<u>A091656</u>	<u>002</u>	Sep 18, 2014
<u>AB</u>			<u>400MG</u>	<u>A091656</u>	<u>003</u>	Sep 18, 2014
<u>AB</u>		DR REDDYS LABS LTD	<u>200MG</u>	<u>A077098</u>	<u>001</u>	Feb 10, 2006
<u>AB</u>			<u>300MG</u>	<u>A077098</u>	<u>002</u>	Feb 10, 2006
<u>AB</u>			<u>400MG</u>	<u>A077098</u>	<u>003</u>	Feb 10, 2006
<u>AB</u>		TEVA	<u>200MG</u>	<u>A076182</u>	<u>001</u>	Sep 02, 2003
<u>AB</u>			<u>300MG</u>	<u>A076182</u>	<u>002</u>	Sep 02, 2003
<u>AB</u>	!		<u>400MG</u>	<u>A076182</u>	<u>003</u>	Sep 02, 2003

OLANZAPINE

INJECTABLE;INTRAMUSCULAR

OLANZAPINE

<u>AP</u>		AM REGENT	<u>10MG/VIAL</u>	<u>A201741</u>	<u>001</u>	Mar 20, 2012
<u>AP</u>		SANDOZ INC	<u>10MG/VIAL</u>	<u>A201588</u>	<u>001</u>	Oct 24, 2011

ZYPREXA

<u>AP</u>	+	LILLY	<u>10MG/VIAL</u>	<u>N021253</u>	<u>001</u>	Mar 29, 2004
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TABLET;ORAL

OLANZAPINE

<u>AB</u>		ALKEM LABS LTD	<u>2.5MG</u>	<u>A202295</u>	<u>001</u>	Oct 20, 2015
<u>AB</u>			<u>5MG</u>	<u>A202295</u>	<u>002</u>	Oct 20, 2015
<u>AB</u>			<u>7.5MG</u>	<u>A202295</u>	<u>003</u>	Oct 20, 2015
<u>AB</u>			<u>10MG</u>	<u>A202295</u>	<u>004</u>	Oct 20, 2015
<u>AB</u>			<u>15MG</u>	<u>A202295</u>	<u>005</u>	Oct 20, 2015
<u>AB</u>			<u>20MG</u>	<u>A202295</u>	<u>006</u>	Oct 20, 2015
<u>AB</u>		APOTEX INC	<u>2.5MG</u>	<u>A090798</u>	<u>001</u>	Apr 23, 2012
<u>AB</u>			<u>5MG</u>	<u>A090798</u>	<u>002</u>	Apr 23, 2012
<u>AB</u>			<u>7.5MG</u>	<u>A090798</u>	<u>003</u>	Apr 23, 2012
<u>AB</u>			<u>10MG</u>	<u>A090798</u>	<u>004</u>	Apr 23, 2012
<u>AB</u>			<u>15MG</u>	<u>A090798</u>	<u>005</u>	Apr 23, 2012
<u>AB</u>			<u>20MG</u>	<u>A090798</u>	<u>006</u>	Apr 23, 2012
<u>AB</u>		AUROBINDO PHARMA LTD	<u>2.5MG</u>	<u>A202050</u>	<u>001</u>	Apr 23, 2012
<u>AB</u>			<u>5MG</u>	<u>A202050</u>	<u>002</u>	Apr 23, 2012
<u>AB</u>			<u>7.5MG</u>	<u>A202050</u>	<u>003</u>	Apr 23, 2012
<u>AB</u>			<u>10MG</u>	<u>A202050</u>	<u>004</u>	Apr 23, 2012
<u>AB</u>			<u>15MG</u>	<u>A202050</u>	<u>005</u>	Apr 23, 2012
<u>AB</u>			<u>20MG</u>	<u>A202050</u>	<u>006</u>	Apr 23, 2012
<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>		HIKMA PHARMS	<u>2.5MG</u>	<u>A204866</u>	<u>001</u>	Jun 16, 2017
<u>AB</u>			<u>5MG</u>	<u>A204866</u>	<u>002</u>	Jun 16, 2017
<u>AB</u>			<u>7.5MG</u>	<u>A204866</u>	<u>003</u>	Jun 16, 2017
<u>AB</u>			<u>10MG</u>	<u>A204866</u>	<u>004</u>	Jun 16, 2017
<u>AB</u>			<u>15MG</u>	<u>A204866</u>	<u>005</u>	Jun 16, 2017
<u>AB</u>			<u>20MG</u>	<u>A204866</u>	<u>006</u>	Jun 16, 2017
<u>AB</u>		INVAGEN PHARMS	<u>2.5MG</u>	<u>A203333</u>	<u>001</u>	Mar 15, 2016
<u>AB</u>			<u>5MG</u>	<u>A203333</u>	<u>002</u>	Mar 15, 2016
<u>AB</u>			<u>7.5MG</u>	<u>A203333</u>	<u>003</u>	Mar 15, 2016
<u>AB</u>			<u>10MG</u>	<u>A203333</u>	<u>004</u>	Mar 15, 2016
<u>AB</u>			<u>15MG</u>	<u>A203333</u>	<u>005</u>	Mar 15, 2016
<u>AB</u>			<u>20MG</u>	<u>A203333</u>	<u>006</u>	Mar 15, 2016
<u>AB</u>		IVAX PHARMS INC	<u>20MG</u>	<u>A077301</u>	<u>001</u>	Apr 29, 2015
<u>AB</u>		JIANGSU HANSON PHARM	<u>2.5MG</u>	<u>A209399</u>	<u>001</u>	Sep 24, 2018
<u>AB</u>			<u>5MG</u>	<u>A209399</u>	<u>002</u>	Sep 24, 2018
<u>AB</u>			<u>10MG</u>	<u>A209399</u>	<u>003</u>	Sep 24, 2018
<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

PRESCRIPTION DRUG PRODUCT LIST

OLANZAPINE

TABLET; ORAL

OLANZAPINE

<u>AB</u>		<u>20MG</u>	<u>A202862</u>	<u>006</u>	Aug 15, 2014
<u>AB</u>	ORCHID HLTHCARE	<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>	SUN PHARM INDS	<u>2.5MG</u>	<u>A091038</u>	<u>001</u>	Apr 23, 2012
<u>AB</u>		<u>5MG</u>	<u>A091038</u>	<u>002</u>	Apr 23, 2012
<u>AB</u>		<u>7.5MG</u>	<u>A091038</u>	<u>003</u>	Apr 23, 2012
<u>AB</u>		<u>10MG</u>	<u>A091038</u>	<u>004</u>	Apr 23, 2012
<u>AB</u>		<u>15MG</u>	<u>A091038</u>	<u>005</u>	Apr 23, 2012
<u>AB</u>		<u>20MG</u>	<u>A091038</u>	<u>006</u>	Apr 23, 2012
<u>AB</u>	TORRENT PHARMS LTD	<u>2.5MG</u>	<u>A091434</u>	<u>001</u>	Apr 23, 2012
<u>AB</u>		<u>5MG</u>	<u>A091434</u>	<u>002</u>	Apr 23, 2012
<u>AB</u>		<u>7.5MG</u>	<u>A091434</u>	<u>003</u>	Apr 23, 2012
<u>AB</u>		<u>10MG</u>	<u>A091434</u>	<u>004</u>	Apr 23, 2012
<u>AB</u>		<u>15MG</u>	<u>A091434</u>	<u>005</u>	Apr 23, 2012
<u>AB</u>		<u>20MG</u>	<u>A091434</u>	<u>006</u>	Apr 23, 2012
<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>	+	LILLY	<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>	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>	INVAGEN PHARMS	<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>	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

PRESCRIPTION DRUG PRODUCT LIST

OLANZAPINE

TABLET, ORALLY DISINTEGRATING;ORAL

OLANZAPINE

<u>AB</u>		<u>10MG</u>	<u>A202285 002</u>	May 12, 2014
<u>AB</u>		<u>15MG</u>	<u>A202285 003</u>	May 12, 2014
<u>AB</u>		<u>20MG</u>	<u>A202285 004</u>	May 12, 2014
<u>AB</u>	ORCHID HLTHCARE	<u>5MG</u>	<u>A202937 001</u>	Mar 02, 2015
<u>AB</u>		<u>10MG</u>	<u>A202937 002</u>	Mar 02, 2015
<u>AB</u>		<u>15MG</u>	<u>A202937 003</u>	Mar 02, 2015
<u>AB</u>		<u>20MG</u>	<u>A202937 004</u>	Mar 02, 2015
<u>AB</u>	PAR PHARM	<u>5MG</u>	<u>A078109 001</u>	Oct 24, 2011
<u>AB</u>		<u>10MG</u>	<u>A078109 002</u>	Oct 24, 2011
<u>AB</u>		<u>15MG</u>	<u>A078109 003</u>	Oct 24, 2011
<u>AB</u>		<u>20MG</u>	<u>A078109 004</u>	Oct 24, 2011
<u>AB</u>	SUN PHARM INDS	<u>5MG</u>	<u>A090881 001</u>	Feb 28, 2012
<u>AB</u>		<u>10MG</u>	<u>A090881 002</u>	Feb 28, 2012
<u>AB</u>		<u>15MG</u>	<u>A090881 003</u>	Feb 28, 2012
<u>AB</u>		<u>20MG</u>	<u>A090881 004</u>	Feb 28, 2012
<u>AB</u>	TORRENT PHARMS LLC	<u>5MG</u>	<u>A091415 001</u>	Oct 25, 2011
<u>AB</u>		<u>10MG</u>	<u>A091415 002</u>	Oct 25, 2011
<u>AB</u>		<u>15MG</u>	<u>A091415 003</u>	Oct 25, 2011
<u>AB</u>		<u>20MG</u>	<u>A091415 004</u>	Oct 25, 2011

ZYPREXA ZYDIS

<u>AB</u>	+!	LILLY	<u>5MG</u>	<u>N021086 001</u>	Apr 06, 2000
<u>AB</u>	+		<u>10MG</u>	<u>N021086 002</u>	Apr 06, 2000
<u>AB</u>	+		<u>15MG</u>	<u>N021086 003</u>	Apr 06, 2000
<u>AB</u>	+		<u>20MG</u>	<u>N021086 004</u>	Apr 06, 2000

OLANZAPINE PAMOATE

SUSPENSION, EXTENDED RELEASE;INTRAMUSCULAR

ZYPREXA RELPREVV

+	ELI LILLY CO	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

OLAPARIB

CAPSULE;ORAL

LYNPARZA

+!	ASTRAZENECA PHARMS	50MG	N206162 001	Dec 19, 2014
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TABLET;ORAL

LYNPARZA

+	ASTRAZENECA PHARMS	100MG	N208558 001	Aug 17, 2017
+!		150MG	N208558 002	Aug 17, 2017

OLIVE OIL; SOYBEAN OIL

EMULSION;INTRAVENOUS

CLINOLIPID 20%

+!	BAXTER HLTHCARE CORP	16%(160GM/1000ML);4% (40GM/1000ML)	N204508 001	Oct 03, 2013
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OLMESARTAN MEDOXOMIL

TABLET;ORAL

BENICAR

<u>AB</u>	+	DAIICHI SANKYO	<u>5MG</u>	<u>N021286 001</u>	Apr 25, 2002
<u>AB</u>	+		<u>20MG</u>	<u>N021286 003</u>	Apr 25, 2002
<u>AB</u>	+!		<u>40MG</u>	<u>N021286 004</u>	Apr 25, 2002

OLMESARTAN MEDOXOMIL

<u>AB</u>		ACCORD HLTHCARE	<u>5MG</u>	<u>A207662 001</u>	Apr 24, 2017
<u>AB</u>			<u>20MG</u>	<u>A207662 002</u>	Apr 24, 2017
<u>AB</u>			<u>40MG</u>	<u>A207662 003</u>	Apr 24, 2017
<u>AB</u>		ALEMBIC PHARMS LTD	<u>5MG</u>	<u>A203012 001</u>	Apr 24, 2017
<u>AB</u>			<u>20MG</u>	<u>A203012 002</u>	Apr 24, 2017
<u>AB</u>			<u>40MG</u>	<u>A203012 003</u>	Apr 24, 2017
<u>AB</u>		ALKEM LABS LTD	<u>5MG</u>	<u>A206763 001</u>	Apr 24, 2017
<u>AB</u>			<u>20MG</u>	<u>A206763 002</u>	Apr 24, 2017
<u>AB</u>			<u>40MG</u>	<u>A206763 003</u>	Apr 24, 2017
<u>AB</u>		AUROBINDO PHARMA LTD	<u>5MG</u>	<u>A204798 001</u>	Apr 24, 2017
<u>AB</u>			<u>20MG</u>	<u>A204798 002</u>	Apr 24, 2017
<u>AB</u>			<u>40MG</u>	<u>A204798 003</u>	Apr 24, 2017
<u>AB</u>		GLENMARK PHARMS LTD	<u>5MG</u>	<u>A203281 001</u>	May 25, 2017
<u>AB</u>			<u>20MG</u>	<u>A203281 002</u>	May 25, 2017
<u>AB</u>			<u>40MG</u>	<u>A203281 003</u>	May 25, 2017
<u>AB</u>		LUPIN LTD	<u>5MG</u>	<u>A206631 001</u>	Apr 27, 2017
<u>AB</u>			<u>20MG</u>	<u>A206631 002</u>	Apr 27, 2017

PRESCRIPTION DRUG PRODUCT LIST

OLMESARTAN MEDOXOMIL

TABLET; ORAL

OLMESARTAN MEDOXOMIL

<u>AB</u>		<u>40MG</u>	<u>A206631 003</u>	Apr 27, 2017
<u>AB</u>	MACLEODS PHARMS LTD	<u>5MG</u>	<u>A204814 001</u>	Apr 24, 2017
<u>AB</u>		<u>20MG</u>	<u>A204814 002</u>	Apr 24, 2017
<u>AB</u>		<u>40MG</u>	<u>A204814 003</u>	Apr 24, 2017
<u>AB</u>	MICRO LABS	<u>5MG</u>	<u>A206372 001</u>	Sep 17, 2019
<u>AB</u>		<u>20MG</u>	<u>A206372 002</u>	Sep 17, 2019
<u>AB</u>		<u>40MG</u>	<u>A206372 003</u>	Sep 17, 2019
<u>AB</u>	MYLAN	<u>5MG</u>	<u>A078276 001</u>	Oct 26, 2016
<u>AB</u>		<u>20MG</u>	<u>A078276 002</u>	Oct 26, 2016
<u>AB</u>		<u>40MG</u>	<u>A078276 003</u>	Oct 26, 2016
<u>AB</u>	QILU	<u>5MG</u>	<u>A210552 001</u>	Jan 10, 2019
<u>AB</u>		<u>20MG</u>	<u>A210552 002</u>	Jan 10, 2019
<u>AB</u>		<u>40MG</u>	<u>A210552 003</u>	Jan 10, 2019
<u>AB</u>	SCIEGEN PHARMS INC	<u>5MG</u>	<u>A208130 001</u>	Jun 29, 2018
<u>AB</u>		<u>20MG</u>	<u>A208130 002</u>	Jun 29, 2018
<u>AB</u>		<u>40MG</u>	<u>A208130 003</u>	Jun 29, 2018
<u>AB</u>	SUNSHINE LAKE	<u>5MG</u>	<u>A211049 001</u>	Feb 22, 2019
<u>AB</u>		<u>20MG</u>	<u>A211049 002</u>	Feb 22, 2019
<u>AB</u>		<u>40MG</u>	<u>A211049 003</u>	Feb 22, 2019
<u>AB</u>	TORRENT	<u>5MG</u>	<u>A202375 001</u>	Apr 24, 2017
<u>AB</u>		<u>20MG</u>	<u>A202375 002</u>	Apr 24, 2017
<u>AB</u>		<u>40MG</u>	<u>A202375 003</u>	Apr 24, 2017
<u>AB</u>	UMEDICA LABS PVT LTD	<u>5MG</u>	<u>A207135 001</u>	Jul 18, 2019
<u>AB</u>		<u>20MG</u>	<u>A207135 002</u>	Jul 18, 2019
<u>AB</u>		<u>40MG</u>	<u>A207135 003</u>	Jul 18, 2019
<u>AB</u>	ZYDUS PHARMS	<u>5MG</u>	<u>A205192 001</u>	Apr 24, 2017
<u>AB</u>		<u>20MG</u>	<u>A205192 002</u>	Apr 24, 2017
<u>AB</u>		<u>40MG</u>	<u>A205192 003</u>	Apr 24, 2017

OLODATEROL HYDROCHLORIDE

SPRAY, METERED; INHALATION

STRIVERDI RESPIMAT

+	BOEHRINGER INGELHEIM	EQ 0.0025MG BASE/INH	N203108 001	Jul 31, 2014
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OLODATEROL HYDROCHLORIDE; TIOTROPIUM BROMIDE

SPRAY, METERED; INHALATION

STIOLTO RESPIMAT

+	BOEHRINGER INGELHEIM	EQ 0.0025MG BASE/INH; EQ 0.0025MG BASE/INH	N206756 001	May 21, 2015
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OLOPATADINE HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

OLOPATADINE HYDROCHLORIDE

<u>AT</u>	AKORN	<u>EQ 0.2% BASE</u>	<u>A204723 001</u>	Dec 05, 2017
<u>AT</u>	AKORN INC	<u>EQ 0.1% BASE</u>	<u>A204532 001</u>	Jan 10, 2017
<u>AT</u>	ALEMBIC PHARMS LTD	<u>EQ 0.1% BASE</u>	<u>A209919 001</u>	Dec 07, 2018
<u>AT</u>		<u>EQ 0.2% BASE</u>	<u>A209420 001</u>	Apr 29, 2019
<u>AT</u>	APOTEX	<u>EQ 0.1% BASE</u>	<u>A078350 001</u>	Dec 07, 2015
<u>AT</u>	APOTEX INC	<u>EQ 0.2% BASE</u>	<u>A090918 001</u>	Dec 05, 2017
<u>AT</u>	AUROBINDO PHARMA LTD	<u>EQ 0.1% BASE</u>	<u>A204812 001</u>	Dec 18, 2015
<u>AT</u>		<u>EQ 0.2% BASE</u>	<u>A209995 001</u>	Apr 04, 2019
<u>AT</u>	BARR LABS INC	<u>EQ 0.2% BASE</u>	<u>A090848 001</u>	Jul 13, 2015
<u>AT</u>	CIPLA	<u>EQ 0.1% BASE</u>	<u>A206046 001</u>	Jul 26, 2017
<u>AT</u>		<u>EQ 0.2% BASE</u>	<u>A206087 001</u>	Dec 05, 2017
<u>AT</u>	FDC LTD	<u>EQ 0.1% BASE</u>	<u>A209282 001</u>	Sep 26, 2019
<u>AT</u>	GLAND PHARMA LTD	<u>EQ 0.1% BASE</u>	<u>A209619 001</u>	Aug 02, 2019
<u>AT</u>	MYLAN	<u>EQ 0.1% BASE</u>	<u>A204392 001</u>	Mar 21, 2018
<u>AT</u>	SOMERSET THERAPS LLC	<u>EQ 0.1% BASE</u>	<u>A206306 001</u>	Dec 07, 2015
<u>AT</u>	USV	<u>EQ 0.1% BASE</u>	<u>A203152 001</u>	Dec 07, 2015
<u>AT</u>	WATSON LABS INC	<u>EQ 0.7% BASE</u>	<u>A208637 001</u>	Feb 19, 2020
<u>AT</u>	WOCKHARDT LTD	<u>EQ 0.1% BASE</u>	<u>A200810 001</u>	Jun 28, 2017
<u>AT</u>	<u>PATADAY</u>			
+	NOVARTIS	<u>EQ 0.2% BASE</u>	<u>N021545 001</u>	Dec 22, 2004
<u>AT</u>	<u>PATANOL</u>			
+	NOVARTIS	<u>EQ 0.1% BASE</u>	<u>N020688 001</u>	Dec 18, 1996
<u>AT</u>	<u>PAZEO</u>			
+	NOVARTIS	<u>EQ 0.7% BASE</u>	<u>N206276 001</u>	Jan 30, 2015

PRESCRIPTION DRUG PRODUCT LIST

OLOPATADINE HYDROCHLORIDE

SPRAY, METERED;NASAL

OLOPATADINE HYDROCHLORIDE

AB	AMNEAL PHARMS LLC	<u>0.665MG/SPRAY</u>	<u>A210901 001</u>	Jan 28, 2020
AB	APOTEX INC	<u>0.665MG/SPRAY</u>	<u>A091572 001</u>	Oct 08, 2014
AB	PERRIGO ISRAEL	<u>0.665MG/SPRAY</u>	<u>A202853 001</u>	Jan 31, 2017

PATANASE

AB	+! NOVARTIS	<u>0.665MG/SPRAY</u>	<u>N021861 001</u>	Apr 15, 2008
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OLSALAZINE SODIUM

CAPSULE;ORAL

DIPENTUM

+!	MYLAN SPECIALITY LP	250MG	N019715 001	Jul 31, 1990
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OMACETAXINE MEPESUCCINATE

POWDER;SUBCUTANEOUS

SYNRIBO

+!	TEVA PHARMS INTL	3.5MG/VIAL	N203585 001	Oct 26, 2012
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OMADACYCLINE TOSYLATE

POWDER;INTRAVENOUS

NUZYRA

+!	PARATEK PHARMS INC	EQ 100MG BASE/VIAL	N209817 001	Oct 02, 2018
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TABLET;ORAL

NUZYRA

+!	PARATEK PHARMS INC	EQ 150MG BASE	N209816 001	Oct 02, 2018
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OMEGA-3-ACID ETHYL ESTERS

CAPSULE;ORAL

LOVAZA

AB	+! SMITHKLINE BEECHAM	<u>1GM CONTAINS AT LEAST 900MG OF THE ETHYL ESTERS OF OMEGA-3 FATTY ACIDS</u>	<u>N021654 001</u>	Nov 10, 2004
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OMEGA-3-ACID ETHYL ESTERS

AB	AMNEAL PHARMS	<u>1GM CONTAINS AT LEAST 900MG OF THE ETHYL ESTERS OF OMEGA-3 FATTY ACIDS</u>	<u>A204940 001</u>	Nov 27, 2015
AB	APOTEX	<u>1GM CONTAINS AT LEAST 900MG OF THE ETHYL ESTERS OF OMEGA-3 FATTY ACIDS</u>	<u>A090973 001</u>	Sep 30, 2014
AB	ASCENT PHARMS INC	<u>1GM CONTAINS AT LEAST 900MG OF THE ETHYL ESTERS OF OMEGA-3 FATTY ACIDS</u>	<u>A207420 001</u>	Feb 25, 2019
AB	BIONPHARMA INC	<u>1GM CONTAINS AT LEAST 900MG OF THE ETHYL ESTERS OF OMEGA-3 FATTY ACIDS</u>	<u>A206455 001</u>	Aug 07, 2019
AB	SOFGEN PHARMS	<u>1GM CONTAINS AT LEAST 900MG OF THE ETHYL ESTERS OF OMEGA-3 FATTY ACIDS</u>	<u>A211355 001</u>	Jul 10, 2019
AB	STRIDES PHARMA	<u>1GM CONTAINS AT LEAST 900MG OF THE ETHYL ESTERS OF OMEGA-3 FATTY ACIDS</u>	<u>A203893 001</u>	Sep 19, 2017
AB	SUN PHARM	<u>1GM CONTAINS AT LEAST 900MG OF THE ETHYL ESTERS OF OMEGA-3 FATTY ACIDS</u>	<u>A210834 001</u>	Jan 09, 2020
AB	TEVA PHARMS USA	<u>1GM CONTAINS AT LEAST 900MG OF THE ETHYL ESTERS OF OMEGA-3 FATTY ACIDS</u>	<u>A091028 001</u>	Apr 07, 2014

OMEPRAZOLE

CAPSULE, DELAYED REL PELLETS;ORAL

OMEPRAZOLE

AB	ACTAVIS LABS FL INC	<u>10MG</u>	<u>A075347 001</u>	May 30, 2008
AB		<u>20MG</u>	<u>A075347 002</u>	May 30, 2008
AB		<u>40MG</u>	<u>A075347 003</u>	May 30, 2008
AB	APOTEX	<u>10MG</u>	<u>A076048 001</u>	Oct 22, 2007
AB		<u>20MG</u>	<u>A076048 002</u>	Oct 22, 2007
AB		<u>40MG</u>	<u>A076048 003</u>	Jan 21, 2009
AB	AUROBINDO PHARMA LTD	<u>10MG</u>	<u>A203270 001</u>	Aug 19, 2015
AB		<u>20MG</u>	<u>A203270 002</u>	Aug 19, 2015
AB		<u>40MG</u>	<u>A203270 003</u>	Aug 19, 2015
AB	BRECKENRIDGE	<u>10MG</u>	<u>A203481 001</u>	Jul 03, 2017
AB		<u>20MG</u>	<u>A203481 002</u>	Jul 03, 2017
AB		<u>40MG</u>	<u>A203481 003</u>	Jul 03, 2017
AB	DR REDDYS LABS LTD	<u>10MG</u>	<u>A075576 003</u>	Oct 22, 2007
AB		<u>10MG</u>	<u>A078490 002</u>	Mar 16, 2009
AB		<u>20MG</u>	<u>A075576 002</u>	Oct 22, 2007
AB		<u>20MG</u>	<u>A078490 003</u>	Mar 16, 2009
AB		<u>40MG</u>	<u>A075576 001</u>	Jan 21, 2009
AB		<u>40MG</u>	<u>A078490 001</u>	Apr 17, 2009
AB	GLENMARK GENERICS	<u>10MG</u>	<u>A091672 001</u>	Oct 31, 2014
AB		<u>20MG</u>	<u>A091672 002</u>	Oct 31, 2014
AB		<u>40MG</u>	<u>A091672 003</u>	Oct 31, 2014
AB	HETERO LABS LTD III	<u>10MG</u>	<u>A204012 001</u>	Sep 26, 2019
AB		<u>20MG</u>	<u>A204012 002</u>	Sep 26, 2019

PRESCRIPTION DRUG PRODUCT LIST

OMEPRAZOLE

CAPSULE, DELAYED REL PELLETS;ORAL

OMEPRAZOLE

<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>		<u>40MG</u>	<u>A204661 002</u>	Jun 13, 2017
<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 PHARMA BV	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>	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 LABS LTD	<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

ZEGERID

<u>AB</u>	+	SALIX	<u>20MG;1.1GM</u>	<u>N021849 001</u>	Feb 27, 2006
<u>AB</u>	+	!	<u>40MG;1.1GM</u>	<u>N021849 002</u>	Feb 27, 2006

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>	PAR PHARM	<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

ZEGERID

<u>AB</u>	+	SALIX	<u>20MG/PACKET;1.68GM/PACKET</u>	<u>N021636 001</u>	Jun 15, 2004
<u>AB</u>	+	!	<u>40MG/PACKET;1.68GM/PACKET</u>	<u>N021636 002</u>	Dec 21, 2004

ONDANSETRON

FILM;ORAL

ZUPLENZ

+	MIDATECH PHARMA US	4MG	N022524 001	Jul 02, 2010
+	!	8MG	N022524 002	Jul 02, 2010

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>	BARR	<u>4MG</u>	<u>A076693 001</u>	Jun 25, 2007
<u>AB</u>		<u>8MG</u>	<u>A076693 002</u>	Jun 25, 2007
<u>AB</u>	GLENMARK GENERICS	<u>4MG</u>	<u>A078152 001</u>	Jun 27, 2007
<u>AB</u>		<u>8MG</u>	<u>A078152 002</u>	Jun 27, 2007
<u>AB</u>	MYLAN	<u>4MG</u>	<u>A078139 001</u>	Jun 25, 2007
<u>AB</u>		<u>8MG</u>	<u>A078139 002</u>	Jun 25, 2007
<u>AB</u>	SANDOZ	<u>4MG</u>	<u>A078050 001</u>	Aug 13, 2007
<u>AB</u>		<u>8MG</u>	<u>A078050 002</u>	Aug 13, 2007
<u>AB</u>	SUN PHARM INDS	<u>4MG</u>	<u>A077557 001</u>	Aug 02, 2007
<u>AB</u>		<u>8MG</u>	<u>A077557 002</u>	Aug 02, 2007
<u>AB</u>	SUN PHARM INDS LTD	<u>4MG</u>	<u>A078602 001</u>	Feb 24, 2011
<u>AB</u>		<u>8MG</u>	<u>A078602 002</u>	Feb 24, 2011
<u>AB</u>	TEVA	<u>4MG</u>	<u>A076810 001</u>	Jun 25, 2007
<u>AB</u>		<u>8MG</u>	<u>A076810 002</u>	Jun 25, 2007

PRESCRIPTION DRUG PRODUCT LIST

ONDANSETRON

TABLET, ORALLY DISINTEGRATING;ORAL

ZOFRAN ODT

<u>AB</u>	+	NOVARTIS	<u>4MG</u>	<u>N020781</u>	<u>001</u>	Jan 27, 1999
<u>AB</u>	+	!	<u>8MG</u>	<u>N020781</u>	<u>002</u>	Jan 27, 1999

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>		AUROBINDO PHARMA LTD	<u>EQ 2MG BASE/ML</u>	<u>A202599</u>	<u>001</u>	Dec 21, 2012
<u>AP</u>	!	BAXTER HLTHCARE CORP	<u>EQ 2MG BASE/ML</u>	<u>A078288</u>	<u>001</u>	Feb 22, 2013
<u>AP</u>		EMCURE PHARMS	<u>EQ 2MG BASE/ML</u>	<u>A090424</u>	<u>001</u>	Apr 16, 2010
<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 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>A077840</u>	<u>001</u>	Jan 19, 2007
<u>AP</u>		MYLAN LABS LTD	<u>EQ 2MG BASE/ML</u>	<u>A204906</u>	<u>001</u>	Jul 31, 2017
<u>AP</u>		QILU	<u>EQ 2MG BASE/ML</u>	<u>A203711</u>	<u>001</u>	Sep 08, 2014
<u>AP</u>		SANDOZ INC	<u>EQ 2MG BASE/ML</u>	<u>A077430</u>	<u>001</u>	Jun 27, 2007
<u>AP</u>		TEVA	<u>EQ 2MG BASE/ML</u>	<u>A076876</u>	<u>001</u>	Nov 22, 2006
<u>AP</u>		WEST-WARD PHARMS INT	<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>		WOCKHARDT	<u>EQ 2MG BASE/ML</u>	<u>A077577</u>	<u>001</u>	Dec 26, 2006

ONDANSETRON HYDROCHLORIDE PRESERVATIVE FREE

<u>AP</u>		ACCORD HLTHCARE	<u>EQ 2MG BASE/ML</u>	<u>A206845</u>	<u>001</u>	Mar 10, 2016
<u>AP</u>		AUROBINDO PHARMA LTD	<u>EQ 2MG BASE/ML</u>	<u>A202600</u>	<u>001</u>	Dec 21, 2012
<u>AP</u>	!	BAXTER HLTHCARE CORP	<u>EQ 2MG BASE/ML</u>	<u>A078287</u>	<u>001</u>	Feb 22, 2013
<u>AP</u>		EMCURE PHARMS LTD	<u>EQ 2MG BASE/ML</u>	<u>A078945</u>	<u>001</u>	Jan 03, 2013
<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>		HOSPIRA	<u>EQ 2MG BASE/ML</u>	<u>A077548</u>	<u>001</u>	Dec 26, 2006
<u>AP</u>		SANDOZ INC	<u>EQ 2MG BASE/ML</u>	<u>A077551</u>	<u>001</u>	Jun 27, 2007
<u>AP</u>		WEST-WARD PHARMS INT	<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>		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>		HIKMA	<u>EQ 4MG BASE/5ML</u>	<u>A076960</u>	<u>001</u>	Dec 26, 2006
<u>AA</u>		LANNETT CO INC	<u>EQ 4MG BASE/5ML</u>	<u>A091342</u>	<u>001</u>	Jan 27, 2011
<u>AA</u>		PHARM ASSOC	<u>EQ 4MG BASE/5ML</u>	<u>A078127</u>	<u>001</u>	Jun 25, 2007
<u>AA</u>		TARO	<u>EQ 4MG BASE/5ML</u>	<u>A077009</u>	<u>001</u>	Nov 30, 2007

ZOFRAN

<u>AA</u>	+	!	NOVARTIS	<u>EQ 4MG BASE/5ML</u>	<u>N020605</u>	<u>001</u>	Jan 24, 1997
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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>			<u>EQ 24MG BASE</u>	<u>A078539</u>	<u>003</u>	Jul 31, 2007
<u>AB</u>		CASI PHARMS INC	<u>EQ 4MG BASE</u>	<u>A077517</u>	<u>001</u>	Jun 25, 2007
<u>AB</u>			<u>EQ 8MG BASE</u>	<u>A077517</u>	<u>002</u>	Jun 25, 2007
<u>AB</u>			<u>EQ 24MG BASE</u>	<u>A077517</u>	<u>003</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>			<u>EQ 24MG BASE</u>	<u>A076183</u>	<u>001</u>	Dec 26, 2006
<u>AB</u>		GLENMARK GENERICS	<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>			<u>EQ 24MG BASE</u>	<u>A077535</u>	<u>003</u>	Jun 25, 2007
<u>AB</u>		IPCA LABS LTD	<u>EQ 4MG BASE</u>	<u>A203761</u>	<u>001</u>	Jan 23, 2014
<u>AB</u>			<u>EQ 8MG BASE</u>	<u>A203761</u>	<u>002</u>	Jan 23, 2014
<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

PRESCRIPTION DRUG PRODUCT LIST

ONDANSETRON HYDROCHLORIDE

TABLET; ORAL

ONDANSETRON HYDROCHLORIDE

<u>AB</u>	PLIVA HRVATSKA DOO	<u>EQ 4MG BASE</u>	<u>A077112 001</u>	Jun 25, 2007
<u>AB</u>		<u>EQ 8MG BASE</u>	<u>A077112 002</u>	Jun 25, 2007
<u>AB</u>		<u>EQ 24MG BASE</u>	<u>A077112 003</u>	Jun 25, 2007
<u>AB</u>	SUN PHARM INDS (IN)	<u>EQ 4MG BASE</u>	<u>A077050 001</u>	Jun 25, 2007
<u>AB</u>		<u>EQ 8MG BASE</u>	<u>A077050 002</u>	Jun 25, 2007
<u>AB</u>	TEVA	<u>EQ 4MG BASE</u>	<u>A076252 001</u>	Jun 25, 2007
<u>AB</u>		<u>EQ 8MG BASE</u>	<u>A076252 002</u>	Jun 25, 2007
<u>AB</u>		<u>EQ 24MG BASE</u>	<u>A076252 003</u>	Jun 25, 2007

ZOFRAN

<u>AB</u>	+ NOVARTIS	<u>EQ 4MG BASE</u>	<u>N020103 001</u>	Dec 31, 1992
<u>AB</u>	+	<u>EQ 8MG BASE</u>	<u>N020103 002</u>	Dec 31, 1992
<u>AB</u>	+!	<u>EQ 24MG BASE</u>	<u>N020103 003</u>	Aug 27, 1999
	ONDANSETRON HYDROCHLORIDE			
	DR REDDYS LABS LTD	EQ 16MG BASE	A076183 004	Dec 26, 2006

ORITAVANCIN DIPHOSPHATE

POWDER; INTRAVENOUS

ORBACTIV

	+!	MELINTA THERAP	EQ 400MG BASE/VIAL	N206334 001	Aug 06, 2014
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ORLISTAT

CAPSULE; ORAL

XENICAL

	+!	CHEPLAPHARM	120MG	N020766 001	Apr 23, 1999
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ORPHENADRINE CITRATE

INJECTABLE; INJECTION

ORPHENADRINE CITRATE

<u>AP</u>	!	AKORN	<u>30MG/ML</u>	<u>A040484 001</u>	May 24, 2006
<u>AP</u>		SAGENT PHARMS	<u>30MG/ML</u>	<u>A090585 001</u>	Aug 30, 2011
<u>AP</u>		WATSON LABS	<u>30MG/ML</u>	<u>A084779 001</u>	Mar 15, 1982
<u>AP</u>		WEST-WARD PHARMS	<u>30MG/ML</u>	<u>A040463 001</u>	Mar 04, 2003
		INT			

TABLET, EXTENDED RELEASE; ORAL

ORPHENADRINE CITRATE

<u>AB</u>		ANDA REPOSITORY	<u>100MG</u>	<u>A040249 001</u>	Jan 29, 1999
<u>AB</u>		IMPAX PHARMS	<u>100MG</u>	<u>A040368 001</u>	Jun 23, 2000
<u>AB</u>		INVAGEN PHARMS	<u>100MG</u>	<u>A091158 001</u>	Jul 27, 2012
<u>AB</u>		LUPIN	<u>100MG</u>	<u>A040284 001</u>	Jun 19, 1998
<u>AB</u>	!	SANDOZ	<u>100MG</u>	<u>A040327 001</u>	Feb 15, 2000

OSELTAMIVIR PHOSPHATE

CAPSULE; ORAL

OSELTAMIVIR PHOSPHATE

<u>AB</u>		ALEMBIC PHARMS LTD	<u>EQ 30MG BASE</u>	<u>A211823 001</u>	Jun 24, 2019
<u>AB</u>			<u>EQ 45MG BASE</u>	<u>A211823 002</u>	Jun 24, 2019
<u>AB</u>			<u>EQ 75MG BASE</u>	<u>A211823 003</u>	Jun 24, 2019
<u>AB</u>		AMNEAL PHARMS	<u>EQ 30MG BASE</u>	<u>A209093 001</u>	May 17, 2017
<u>AB</u>			<u>EQ 45MG BASE</u>	<u>A209093 002</u>	May 17, 2017
<u>AB</u>			<u>EQ 75MG BASE</u>	<u>A209093 003</u>	May 17, 2017
<u>AB</u>		HETERO LABS LTD III	<u>EQ 30MG BASE</u>	<u>A209438 001</u>	Feb 23, 2018
<u>AB</u>			<u>EQ 45MG BASE</u>	<u>A209438 002</u>	Feb 23, 2018
<u>AB</u>			<u>EQ 75MG BASE</u>	<u>A209438 003</u>	Feb 23, 2018
<u>AB</u>		LUPIN ATLANTIS	<u>EQ 30MG BASE</u>	<u>A208348 001</u>	Jan 09, 2018
<u>AB</u>			<u>EQ 45MG BASE</u>	<u>A208348 002</u>	Jan 09, 2018
<u>AB</u>			<u>EQ 75MG BASE</u>	<u>A208348 003</u>	Jan 09, 2018
<u>AB</u>		MACLEODS PHARMS LTD	<u>EQ 30MG BASE</u>	<u>A207211 001</u>	Sep 14, 2017
<u>AB</u>			<u>EQ 45MG BASE</u>	<u>A207211 002</u>	Sep 14, 2017
<u>AB</u>			<u>EQ 75MG BASE</u>	<u>A207211 003</u>	Sep 14, 2017
<u>AB</u>		NATCO PHARMA LTD	<u>EQ 30MG BASE</u>	<u>A202595 001</u>	Aug 03, 2016
<u>AB</u>			<u>EQ 45MG BASE</u>	<u>A202595 002</u>	Aug 03, 2016
<u>AB</u>			<u>EQ 75MG BASE</u>	<u>A202595 003</u>	Aug 03, 2016
<u>AB</u>		NESHER PHARMS	<u>EQ 30MG BASE</u>	<u>A208578 001</u>	Feb 24, 2017
<u>AB</u>			<u>EQ 45MG BASE</u>	<u>A208578 002</u>	Feb 24, 2017
<u>AB</u>			<u>EQ 75MG BASE</u>	<u>A208578 003</u>	Feb 24, 2017
<u>AB</u>		STRIDES PHARMA	<u>EQ 30MG BASE</u>	<u>A209421 001</u>	Jun 08, 2018
<u>AB</u>			<u>EQ 45MG BASE</u>	<u>A209421 002</u>	Jun 08, 2018
<u>AB</u>			<u>EQ 75MG BASE</u>	<u>A209421 003</u>	Jun 08, 2018
<u>AB</u>		SUNSHINE LAKE	<u>EQ 30MG BASE</u>	<u>A212739 001</u>	Mar 04, 2020
<u>AB</u>			<u>EQ 45MG BASE</u>	<u>A212739 002</u>	Mar 04, 2020
<u>AB</u>			<u>EQ 75MG BASE</u>	<u>A212739 003</u>	Mar 04, 2020

PRESCRIPTION DRUG PRODUCT LIST

OSELTAMIVIR PHOSPHATE

CAPSULE; ORAL

TAMIFLU

AB	+	ROCHE	<u>EQ 30MG BASE</u>	<u>N021087 003</u>	Jul 02, 2007
AB	+		<u>EQ 45MG BASE</u>	<u>N021087 002</u>	Jul 02, 2007
AB	+	!	<u>EQ 75MG BASE</u>	<u>N021087 001</u>	Oct 27, 1999

FOR SUSPENSION; ORAL

OSELTAMIVIR PHOSPHATE

AB		ALVOGEN PINE BROOK	<u>EQ 6MG BASE/ML</u>	<u>A208823 001</u>	Oct 31, 2017
AB		AMNEAL PHARMS NY	<u>EQ 6MG BASE/ML</u>	<u>A210186 001</u>	Feb 27, 2018
AB		LUPIN ATLANTIS	<u>EQ 6MG BASE/ML</u>	<u>A208347 001</u>	Feb 20, 2018
AB		NESHER PHARMS	<u>EQ 6MG BASE/ML</u>	<u>A209113 001</u>	Sep 14, 2017
AB		TEVA PHARMS USA	<u>EQ 6MG BASE/ML</u>	<u>A211125 001</u>	Feb 27, 2019

TAMIFLU

AB	+	!	ROCHE	<u>EQ 6MG BASE/ML</u>	<u>N021246 002</u>	Mar 21, 2011
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OSIMERTINIB MESYLATE

TABLET; ORAL

TAGRISSO

	+	ASTRAZENECA PHARMS	EQ 40MG BASE	N208065 001	Nov 13, 2015	
	+	!		EQ 80MG BASE	N208065 002	Nov 13, 2015

OSPEMIFENE

TABLET; ORAL

OSPHENA

	+	!	DUCHESNAY	60MG	N203505 001	Feb 26, 2013
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OXACILLIN SODIUM

INJECTABLE; INJECTION

OXACILLIN SODIUM

AP		AUROBINDO PHARMA LTD	<u>EQ 1GM BASE/VIAL</u>	<u>A201539 001</u>	Jan 18, 2013
AP			<u>EQ 2GM BASE/VIAL</u>	<u>A201539 002</u>	Jan 18, 2013
AP			<u>EQ 10GM BASE/VIAL</u>	<u>A201538 001</u>	Jan 18, 2013
AP		PIRAMAL CRITICAL	<u>EQ 1GM BASE/VIAL</u>	<u>A206681 001</u>	Sep 11, 2017
AP			<u>EQ 2GM BASE/VIAL</u>	<u>A206681 002</u>	Sep 11, 2017
AP			<u>EQ 10GM BASE/VIAL</u>	<u>A206760 001</u>	Oct 26, 2017
AP	!	SAGENT PHARMS	<u>EQ 1GM BASE/VIAL</u>	<u>A091246 001</u>	Mar 30, 2012
AP	!		<u>EQ 2GM BASE/VIAL</u>	<u>A091246 002</u>	Mar 30, 2012
AP	!		<u>EQ 10GM BASE/VIAL</u>	<u>A091245 001</u>	Mar 30, 2012
AP		WOCKHARDT BIO AG	<u>EQ 1GM BASE/VIAL</u>	<u>A207147 001</u>	Jul 31, 2017
AP			<u>EQ 2GM BASE/VIAL</u>	<u>A207147 002</u>	Jul 31, 2017
AP			<u>EQ 10GM BASE/VIAL</u>	<u>A207148 001</u>	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

ELOXATIN

AP	+	!	SANOFI AVENTIS US	<u>50MG/10ML (5MG/ML)</u>	<u>N021759 001</u>	Jan 31, 2005
AP	+	!		<u>100MG/20ML (5MG/ML)</u>	<u>N021759 002</u>	Jan 31, 2005

OXALIPLATIN

AP		ACCORD HLTHCARE	<u>50MG/10ML (5MG/ML)</u>	<u>A207474 001</u>	Mar 21, 2017
AP			<u>100MG/20ML (5MG/ML)</u>	<u>A207474 002</u>	Mar 21, 2017
AP			<u>200MG/40ML (5MG/ML)</u>	<u>A207474 003</u>	Mar 21, 2017
AP		ACTAVIS LLC	<u>50MG/10ML (5MG/ML)</u>	<u>A204880 001</u>	Mar 05, 2018
AP			<u>100MG/20ML (5MG/ML)</u>	<u>A204880 002</u>	Mar 05, 2018
AP	!	ACTAVIS TOTOWA	<u>50MG/VIAL</u>	<u>A078803 001</u>	Aug 08, 2012
AP	!		<u>100MG/VIAL</u>	<u>A078803 002</u>	Aug 08, 2012
AP		CIPLA	<u>50MG/10ML (5MG/ML)</u>	<u>A208523 001</u>	Feb 10, 2017
AP			<u>100MG/20ML (5MG/ML)</u>	<u>A208523 002</u>	Feb 10, 2017
AP		EUGIA PHARMA	<u>50MG/10ML (5MG/ML)</u>	<u>A205529 001</u>	Sep 06, 2017
AP			<u>100MG/20ML (5MG/ML)</u>	<u>A205529 002</u>	Sep 06, 2017
AP		FRESENIUS KABI USA	<u>50MG/10ML (5MG/ML)</u>	<u>A078811 001</u>	Jun 10, 2010
AP			<u>100MG/20ML (5MG/ML)</u>	<u>A078811 002</u>	Jun 10, 2010
AP			<u>50MG/VIAL</u>	<u>A078819 001</u>	Jun 02, 2010
AP			<u>50MG/10ML (5MG/ML)</u>	<u>A090030 001</u>	Jan 31, 2017
AP			<u>100MG/VIAL</u>	<u>A078819 002</u>	Jun 02, 2010
AP			<u>100MG/20ML (5MG/ML)</u>	<u>A090030 002</u>	Jan 31, 2017
AP			<u>200MG/40ML (5MG/ML)</u>	<u>A090030 003</u>	Jan 31, 2017
AP		GLAND PHARMA LTD	<u>50MG/10ML (5MG/ML)</u>	<u>A207325 001</u>	Feb 10, 2017
AP			<u>50MG/VIAL</u>	<u>A207385 001</u>	May 23, 2017
AP			<u>100MG/20ML (5MG/ML)</u>	<u>A207325 002</u>	Feb 10, 2017
AP			<u>100MG/VIAL</u>	<u>A207385 002</u>	May 23, 2017

PRESCRIPTION DRUG PRODUCT LIST

OXALIPLATIN

INJECTABLE; INTRAVENOUS

OXALIPLATIN

<u>AP</u>		<u>200MG/40ML (5MG/ML)</u>	<u>A207325 003</u>	Oct 18, 2017
<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>	INGENUS PHARMS LLC	<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>	JIANGSU HENGRUI MED	<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>	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>	QILU	<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>	!	<u>200MG/40ML (5MG/ML)</u>	<u>A204368 003</u>	Jun 07, 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>	+!	<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

OXANDROLONE

TABLET; ORAL

OXANDROLONE

<u>AB</u>	PAR PHARM	<u>2.5MG</u>	<u>A077827 001</u>	Jun 22, 2007
<u>AB</u>	!	<u>10MG</u>	<u>A077827 002</u>	Jun 22, 2007
<u>AB</u>	UPSHER SMITH LABS	<u>2.5MG</u>	<u>A076761 001</u>	Dec 01, 2006
<u>AB</u>		<u>10MG</u>	<u>A078033 001</u>	Mar 22, 2007

OXAPROZIN

TABLET; ORAL

DAYPRO

<u>AB</u>	+!	GD SEARLE	<u>600MG</u>	<u>N018841 004</u>	Oct 29, 1992
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OXAPROZIN

<u>AB</u>	AMNEAL PHARMS CO	<u>600MG</u>	<u>A208633 001</u>	May 04, 2017
<u>AB</u>	APOTEX INC	<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>	IVAX SUB TEVA PHARMS	<u>600MG</u>	<u>A075846 001</u>	May 13, 2002
<u>AB</u>	SANDOZ	<u>600MG</u>	<u>A075845 001</u>	Jan 31, 2001
<u>AB</u>	TEVA	<u>600MG</u>	<u>A075849 001</u>	Jul 03, 2002

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>	SANDOZ	<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

OXCARBAZEPINE

SUSPENSION; ORAL

OXCARBAZEPINE

<u>AB</u>	AMNEAL PHARMS	<u>300MG/5ML</u>	<u>A202961 001</u>	Sep 17, 2012
<u>AB</u>	HIKMA	<u>300MG/5ML</u>	<u>A201193 001</u>	Oct 03, 2012
<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
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TABLET; ORAL

OXCARBAZEPINE

<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>300MG</u>	<u>A077747 002</u>	Apr 09, 2008
<u>AB</u>		<u>600MG</u>	<u>A077747 003</u>	Apr 09, 2008
<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

PRESCRIPTION DRUG PRODUCT LIST

OXCARBAZEPINE

TABLET; ORAL

OXCARBAZEPINE

<u>AB</u>	TARO	<u>150MG</u>	<u>A077801</u>	<u>001</u>	Nov 15, 2007
<u>AB</u>		<u>300MG</u>	<u>A077801</u>	<u>002</u>	Nov 15, 2007
<u>AB</u>		<u>600MG</u>	<u>A077801</u>	<u>003</u>	Nov 15, 2007

TRILEPTAL

<u>AB</u>	+	NOVARTIS	<u>150MG</u>	<u>N021014</u>	<u>001</u>	Jan 14, 2000
<u>AB</u>	+		<u>300MG</u>	<u>N021014</u>	<u>002</u>	Jan 14, 2000
<u>AB</u>	+	!	<u>600MG</u>	<u>N021014</u>	<u>003</u>	Jan 14, 2000

TABLET, EXTENDED RELEASE; ORAL

OXTELLAR XR

+	SUPERNUS PHARMS	150MG	N202810	001	Oct 19, 2012
+		300MG	N202810	002	Oct 19, 2012
+	!	600MG	N202810	003	Oct 19, 2012

OXICONAZOLE NITRATE

CREAM; TOPICAL

OXICONAZOLE NITRATE

<u>AB</u>	TARO PHARMS	<u>EQ 1% BASE</u>	<u>A205076</u>	<u>001</u>	Mar 07, 2016
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OXISTAT

<u>AB</u>	+	FOUGERA PHARMS	<u>EQ 1% BASE</u>	<u>N019828</u>	<u>001</u>	Dec 30, 1988
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LOTION; TOPICAL

OXISTAT

+	!	FOUGERA PHARMS	EQ 1% BASE	N020209	001	Sep 30, 1992
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OXYBUTYNIN

FILM, EXTENDED RELEASE; TRANSDERMAL

OXYTROL

+	!	ALLERGAN	3.9MG/24HR	N021351	002	Feb 26, 2003
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OXYBUTYNIN CHLORIDE

GEL; TRANSDERMAL

GELNIQUE

+	!	ALLERGAN	10% (100MG/PACKET)	N022204	001	Jan 27, 2009
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SYRUP; ORAL

OXYBUTYNIN CHLORIDE

<u>AA</u>	LANNETT CO INC	<u>5MG/5ML</u>	<u>A074520</u>	<u>001</u>	Mar 29, 1996	
<u>AA</u>	PHARM ASSOC	<u>5MG/5ML</u>	<u>A075137</u>	<u>001</u>	Dec 18, 1998	
<u>AA</u>	!	WOCKHARDT BIO AG	<u>5MG/5ML</u>	<u>A074868</u>	<u>001</u>	Feb 12, 1997

TABLET; ORAL

OXYBUTYNIN CHLORIDE

<u>AB</u>	ABHAI LLC	<u>5MG</u>	<u>A209335</u>	<u>001</u>	Dec 22, 2017	
<u>AB</u>	ARISE	<u>5MG</u>	<u>A209025</u>	<u>001</u>	Dec 21, 2017	
<u>AB</u>	ATHEM	<u>5MG</u>	<u>A211062</u>	<u>001</u>	Feb 06, 2019	
<u>AB</u>	EMCURE PHARMS LTD	<u>5MG</u>	<u>A211682</u>	<u>001</u>	May 10, 2019	
<u>AB</u>	NOVAST LABS	<u>5MG</u>	<u>A210611</u>	<u>001</u>	Oct 30, 2019	
<u>AB</u>	NOVITIUM PHARMA	<u>5MG</u>	<u>A209823</u>	<u>001</u>	Oct 23, 2017	
<u>AB</u>	TEVA PHARMS USA	<u>5MG</u>	<u>A071655</u>	<u>001</u>	Nov 14, 1988	
<u>AB</u>	TULEX PHARMS INC	<u>5MG</u>	<u>A210125</u>	<u>001</u>	Sep 06, 2018	
<u>AB</u>	UPSHER SMITH LABS	<u>5MG</u>	<u>A074625</u>	<u>001</u>	Jul 31, 1996	
<u>AB</u>	!	VINTAGE PHARMS	<u>5MG</u>	<u>A075079</u>	<u>001</u>	Oct 31, 1997

TABLET, EXTENDED RELEASE; ORAL

DITROPAN XL

<u>AB</u>	+	JANSSEN PHARMS	<u>5MG</u>	<u>N020897</u>	<u>001</u>	Dec 16, 1998
<u>AB</u>	+		<u>10MG</u>	<u>N020897</u>	<u>002</u>	Dec 16, 1998

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 INC	<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>	IMPAX PHARMS	<u>5MG</u>	<u>A076745</u>	<u>002</u>	May 09, 2007
<u>AB</u>		<u>10MG</u>	<u>A076745</u>	<u>003</u>	May 09, 2007
<u>AB</u>		<u>15MG</u>	<u>A076745</u>	<u>001</u>	Nov 09, 2006
<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

PRESCRIPTION DRUG PRODUCT LIST

OXYBUTYNIN CHLORIDE

TABLET, EXTENDED RELEASE;ORAL

OXYBUTYNIN CHLORIDE

<u>AB</u>		<u>15MG</u>	<u>A078503</u>	<u>003</u>	Feb 04, 2009
<u>AB</u>	UNIQUE PHARM LABS	<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 INC	<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>5MG</u>	<u>N200534</u>	<u>001</u>	Oct 20, 2010
<u>AB</u>	LANNETT CO INC	<u>5MG</u>	<u>A203823</u>	<u>001</u>	Aug 01, 2014
<u>AB</u>	MAYNE PHARMA INC	<u>5MG</u>	<u>A203107</u>	<u>001</u>	Jul 26, 2012
<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 INC	<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>	AUROLIFE PHARMA LLC	<u>5MG/5ML</u>	<u>A212429</u>	<u>001</u>	Jan 27, 2020
<u>AA</u>		<u>100MG/5ML</u>	<u>A212429</u>	<u>002</u>	Jan 27, 2020
<u>AA</u>	EYWA	<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
<u>AA</u>	+ GENUS LIFESCIENCES	<u>5MG/5ML</u>	<u>N200535</u>	<u>002</u>	Aug 22, 2013
<u>AA</u>	+!	<u>100MG/5ML</u>	<u>N200535</u>	<u>001</u>	Oct 20, 2010
<u>AA</u>	HI TECH	<u>5MG/5ML</u>	<u>A208817</u>	<u>001</u>	Aug 10, 2017
<u>AA</u>		<u>100MG/5ML</u>	<u>A208795</u>	<u>001</u>	Aug 07, 2017
<u>AA</u>	HIKMA	<u>5MG/5ML</u>	<u>A204037</u>	<u>001</u>	Jul 15, 2013
<u>AA</u>	MAYNE PHARMA INC	<u>100MG/5ML</u>	<u>A204092</u>	<u>001</u>	Jun 05, 2014
<u>AA</u>	NOVEL LABS INC	<u>100MG/5ML</u>	<u>A204603</u>	<u>001</u>	Apr 29, 2015
<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>	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>	+! VISTAPHARM	<u>5MG/5ML</u>	<u>N201194</u>	<u>001</u>	Jan 12, 2012
<u>AA</u>		<u>100MG/5ML</u>	<u>A202537</u>	<u>001</u>	Jul 30, 2012
<u>AA</u>	WOCKHARDT BIO AG	<u>5MG/5ML</u>	<u>A206456</u>	<u>001</u>	Jun 16, 2015

TABLET;ORAL

OXYCODONE HYDROCHLORIDE

<u>AB</u>	ACTAVIS ELIZABETH	<u>5MG</u>	<u>A076636</u>	<u>003</u>	Apr 07, 2015
<u>AB</u>		<u>15MG</u>	<u>A076636</u>	<u>001</u>	Feb 06, 2004
<u>AB</u>		<u>30MG</u>	<u>A076636</u>	<u>002</u>	Feb 06, 2004
<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
<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

PRESCRIPTION DRUG PRODUCT LIST

OXYCODONE HYDROCHLORIDE

TABLET; ORAL

OXYCODONE HYDROCHLORIDE

<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>	MAYNE PHARMA INC	<u>5MG</u>	<u>A091313</u>	<u>001</u>	Feb 18, 2011
<u>AB</u>		<u>10MG</u>	<u>A091313</u>	<u>004</u>	Apr 29, 2016
<u>AB</u>		<u>15MG</u>	<u>A091313</u>	<u>002</u>	Feb 18, 2011
<u>AB</u>		<u>20MG</u>	<u>A091313</u>	<u>005</u>	Apr 29, 2016
<u>AB</u>		<u>30MG</u>	<u>A091313</u>	<u>003</u>	Feb 18, 2011
<u>AB</u>	NESHER PHARMS	<u>5MG</u>	<u>A077290</u>	<u>001</u>	Dec 08, 2005
<u>AB</u>		<u>10MG</u>	<u>A077290</u>	<u>002</u>	Dec 08, 2005
<u>AB</u>		<u>15MG</u>	<u>A077290</u>	<u>003</u>	Dec 08, 2005
<u>AB</u>		<u>20MG</u>	<u>A077290</u>	<u>004</u>	Dec 08, 2005
<u>AB</u>		<u>30MG</u>	<u>A077290</u>	<u>005</u>	Dec 08, 2005
<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>15MG</u>	<u>A076758</u>	<u>001</u>	Jun 30, 2004
<u>AB</u>		<u>30MG</u>	<u>A076758</u>	<u>002</u>	Jun 30, 2004
<u>AB</u>	SUN PHARM INDS INC	<u>5MG</u>	<u>A090659</u>	<u>001</u>	Apr 10, 2009
<u>AB</u>		<u>10MG</u>	<u>A090659</u>	<u>005</u>	Nov 06, 2012
<u>AB</u>		<u>15MG</u>	<u>A090659</u>	<u>002</u>	Apr 10, 2009
<u>AB</u>		<u>20MG</u>	<u>A090659</u>	<u>004</u>	Nov 06, 2012
<u>AB</u>		<u>30MG</u>	<u>A090659</u>	<u>003</u>	Apr 10, 2009

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

OXAYDO

ZYLA	5MG	N202080	001	Jun 17, 2011
	7.5MG	N202080	002	Jun 17, 2011

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

PRESCRIPTION DRUG PRODUCT LIST

OXYMETAZOLINE HYDROCHLORIDE

CREAM; TOPICAL

RHOFADE

+! EPI HLTH 1% N208552 001 Jan 18, 2017

OXYMETAZOLINE HYDROCHLORIDE; TETRACAINE HYDROCHLORIDE

SPRAY, METERED; NASAL

KOVANAZE

+! ST RENATUS 0.1MG/SPRAY; 6MG/SPRAY N208032 001 Jun 29, 2016

OXYMETHOLONE

TABLET; ORAL

ANADROL-50

+! MYLAN SPECIALITY LP 50MG N016848 001

OXYMORPHONE HYDROCHLORIDE

TABLET; ORAL

OXYMORPHONE HYDROCHLORIDE

<u>AB</u>	ASCENT PHARMS INC	<u>5MG</u>	<u>A210175 001</u>	Feb 02, 2018
<u>AB</u>		<u>10MG</u>	<u>A210175 002</u>	Feb 02, 2018
<u>AB</u>	AUROLIFE PHARMA LLC	<u>5MG</u>	<u>A204459 001</u>	Apr 26, 2016
<u>AB</u>		<u>10MG</u>	<u>A204459 002</u>	Apr 26, 2016
<u>AB</u>	AVANTHI INC	<u>5MG</u>	<u>A203601 001</u>	Jan 30, 2013
<u>AB</u>	!	<u>10MG</u>	<u>A203601 002</u>	Jan 30, 2013
<u>AB</u>	EPIC PHARMA LLC	<u>5MG</u>	<u>A201187 001</u>	Dec 15, 2014
<u>AB</u>		<u>10MG</u>	<u>A201187 002</u>	Dec 15, 2014
<u>AB</u>	HIKMA	<u>5MG</u>	<u>A090964 001</u>	Sep 27, 2010
<u>AB</u>		<u>10MG</u>	<u>A090964 002</u>	Sep 27, 2010
<u>AB</u>	SPECGX LLC	<u>5MG</u>	<u>A202321 001</u>	Apr 25, 2013
<u>AB</u>		<u>10MG</u>	<u>A202321 002</u>	Apr 25, 2013
<u>AB</u>	TEVA	<u>5MG</u>	<u>A091443 002</u>	Feb 15, 2011
<u>AB</u>		<u>10MG</u>	<u>A091443 001</u>	Feb 15, 2011

TABLET, EXTENDED RELEASE; ORAL

OXYMORPHONE HYDROCHLORIDE

<u>AB</u>	ACTAVIS ELIZABETH	<u>5MG</u>	<u>A079046 003</u>	Jul 11, 2013
<u>AB</u>		<u>7.5MG</u>	<u>A079046 001</u>	Dec 13, 2010
<u>AB</u>		<u>10MG</u>	<u>A079046 004</u>	Jul 11, 2013
<u>AB</u>		<u>15MG</u>	<u>A079046 002</u>	Dec 13, 2010
<u>AB</u>		<u>20MG</u>	<u>A079046 005</u>	Jul 11, 2013
<u>AB</u>		<u>30MG</u>	<u>A079046 006</u>	Jul 11, 2013
<u>AB</u>		<u>40MG</u>	<u>A079046 007</u>	Jul 11, 2013
<u>AB</u>	HIKMA	<u>5MG</u>	<u>A200822 002</u>	Jul 15, 2013
<u>AB</u>		<u>7.5MG</u>	<u>A200822 003</u>	Jul 15, 2013
<u>AB</u>		<u>10MG</u>	<u>A200822 004</u>	Jul 15, 2013
<u>AB</u>		<u>15MG</u>	<u>A200822 005</u>	Jul 15, 2013
<u>AB</u>		<u>20MG</u>	<u>A200822 006</u>	Jul 15, 2013
<u>AB</u>		<u>30MG</u>	<u>A200822 007</u>	Jul 15, 2013
<u>AB</u>		<u>40MG</u>	<u>A200822 001</u>	Jul 15, 2013
<u>AB</u>	IMPAX LABS	<u>5MG</u>	<u>A079087 001</u>	Jun 14, 2010
<u>AB</u>		<u>7.5MG</u>	<u>A079087 002</u>	Dec 21, 2010
<u>AB</u>		<u>10MG</u>	<u>A079087 003</u>	Jun 14, 2010
<u>AB</u>		<u>15MG</u>	<u>A079087 004</u>	Dec 21, 2010
<u>AB</u>		<u>20MG</u>	<u>A079087 005</u>	Jun 14, 2010
<u>AB</u>		<u>30MG</u>	<u>A079087 006</u>	Jul 22, 2010
<u>AB</u>	SPECGX LLC	<u>5MG</u>	<u>A202946 001</u>	Jun 27, 2014
<u>AB</u>		<u>7.5MG</u>	<u>A202946 002</u>	Jun 27, 2014
<u>AB</u>		<u>10MG</u>	<u>A202946 003</u>	Jun 27, 2014
<u>AB</u>		<u>15MG</u>	<u>A202946 004</u>	Jun 27, 2014
<u>AB</u>		<u>20MG</u>	<u>A202946 005</u>	Jun 27, 2014
<u>AB</u>		<u>30MG</u>	<u>A202946 006</u>	Jun 27, 2014
<u>AB</u>		<u>40MG</u>	<u>A202946 007</u>	Jun 27, 2014
<u>AB</u>	!	IMPAX LABS	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 001</u>
<u>AP</u>	+!		<u>100USP UNITS/10ML (10USP UNITS/ML)</u>	<u>N018248 002</u>
<u>AP</u>		HIKMA FARMACEUTICA	<u>10USP UNITS/ML (10USP UNITS/ML)</u>	<u>A200219 001</u> Feb 13, 2013
<u>AP</u>		SAGENT PHARMS INC	<u>10USP UNITS/ML (10USP UNITS/ML)</u>	<u>A091676 001</u> Jul 13, 2018
<u>AP</u>			<u>100USP UNITS/10ML (10USP UNITS/ML)</u>	<u>A091676 002</u> Jul 13, 2018
<u>AP</u>	+!	WEST-WARD PHARMS	<u>10USP UNITS/ML (10USP UNITS/ML)</u>	<u>N018243 001</u>
<u>AP</u>	+!	INT	<u>100USP UNITS/10ML (10USP UNITS/ML)</u>	<u>N018243 002</u> Jan 10, 2007

PRESCRIPTION DRUG PRODUCT LIST

OXYTOCIN

INJECTABLE; INJECTION

PITOCIN

AP	+ !	PAR STERILE PRODUCTS	10USP UNITS/ML (10USP UNITS/ML)	N018261 001	
AP	+		100USP UNITS/10ML (10USP UNITS/ML)	N018261 002	Jul 27, 2007
		OXYTOCIN			
	+ !	FRESENIUS KABI USA	300USP UNITS/30ML (10USP UNITS/ML)	N018248 003	Jul 27, 2007
		PITOCIN			
	+	PAR STERILE PRODUCTS	500USP UNITS/50ML (10USP UNITS/ML)	N018261 003	Sep 05, 2012

OZENOXACIN

CREAM; TOPICAL

XEPI

	+ !	FERRER INTERNACIONAL	1%	N208945 001	Dec 11, 2017
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PACLITAXEL

FOR SUSPENSION; IV (INFUSION)

ABRAXANE

	+ !	ABRAXIS BIOSCIENCE	100MG/VIAL	N021660 001	Jan 07, 2005
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INJECTABLE; INJECTION

PACITAXEL

AP		GLAND PHARMA LTD	6MG/ML	A207326 001	Aug 23, 2016
		<u>PACLITAXEL</u>			
AP		ACTAVIS TOTOWA	6MG/ML	A090130 001	Dec 09, 2009
AP		FRESENIUS KABI USA	6MG/ML	A077574 001	Nov 27, 2006
AP	!	HOSPIRA	6MG/ML	A076131 001	May 08, 2002
AP		MYLAN LABS LTD	6MG/ML	A091540 001	Sep 29, 2011
AP		TEVA PHARMS	6MG/ML	A075184 001	Jan 25, 2002
AP		WEST-WARD PHARMS INT	6MG/ML	A075190 001	Jan 28, 2002

PALBOCICLIB

CAPSULE; ORAL

IBRANCE

	+	PFIZER INC	75MG	N207103 001	Feb 03, 2015
	+		100MG	N207103 002	Feb 03, 2015
	+ !		125MG	N207103 003	Feb 03, 2015

TABLET; ORAL

IBRANCE

	+	PFIZER INC	75MG	N212436 001	Nov 01, 2019
	+		100MG	N212436 002	Nov 01, 2019
	+ !		125MG	N212436 003	Nov 01, 2019

PALIPERIDONE

TABLET, EXTENDED RELEASE; ORAL

INVEGA

AB	+	JANSSEN PHARMS	1.5MG	N021999 006	Aug 26, 2008
AB	+		3MG	N021999 001	Dec 19, 2006
AB	+ !		6MG	N021999 002	Dec 19, 2006
AB	+		9MG	N021999 003	Dec 19, 2006

PALIPERIDONE

AB		ACTAVIS LABS FL INC	1.5MG	A202645 001	Aug 03, 2015
AB			3MG	A202645 002	Aug 03, 2015
AB			6MG	A202645 003	Aug 03, 2015
AB			9MG	A202645 004	Aug 03, 2015
AB		AMNEAL PHARMS	1.5MG	A204707 001	Sep 23, 2019
AB			3MG	A204707 002	Sep 23, 2019
AB			6MG	A204707 003	Sep 23, 2019
AB			9MG	A204707 004	Sep 23, 2019
AB		INVENTIA	1.5MG	A204452 001	Jun 12, 2019
AB			3MG	A204452 002	Jun 12, 2019
AB			6MG	A204452 003	Jun 12, 2019
AB			9MG	A204452 004	Jun 12, 2019
AB		MYLAN	1.5MG	A203802 001	Sep 24, 2015
AB			3MG	A203802 002	Sep 24, 2015
AB			6MG	A203802 003	Sep 24, 2015
AB			9MG	A203802 004	Sep 24, 2015
AB		SUN PHARM	1.5mg	A205618 001	Apr 06, 2018
AB			3MG	A205618 002	Apr 06, 2018
AB			6MG	A205618 003	Apr 06, 2018
AB			9MG	A205618 004	Apr 06, 2018

PRESCRIPTION DRUG PRODUCT LIST

PALIPERIDONE PALMITATE

SUSPENSION, EXTENDED RELEASE; INTRAMUSCULAR

INVEGA SUSTENNA

+	JANSSEN PHARMS	39MG/0.25ML (39MG/0.25ML)	N022264	001	Jul 31, 2009
+		78MG/0.5ML (78MG/0.5ML)	N022264	002	Jul 31, 2009
+		117MG/0.75ML (117MG/0.75ML)	N022264	003	Jul 31, 2009
+		156MG/ML (156MG/ML)	N022264	004	Jul 31, 2009
+		234MG/1.5ML (156MG/ML)	N022264	005	Jul 31, 2009

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

ALOXI

AP +! HELSINN HLTHCARE EQ 0.075MG BASE/1.5ML (EQ 0.05MG BASE/ML) N021372 002 Feb 29, 2008

AP +! EQ 0.25MG BASE/5ML (EQ 0.05MG BASE/ML) N021372 001 Jul 25, 2003

PALONOSETRON HYDROCHLORIDE

AP AUROBINDO PHARMA LTD EQ 0.25MG BASE/5ML (EQ 0.05MG BASE/ML) A204702 001 Nov 06, 2018

AP CIPLA EQ 0.25MG BASE/5ML (EQ 0.05MG BASE/ML) A206396 001 Sep 19, 2018

AP DR REDDYS LABS LTD EQ 0.25MG BASE/5ML (EQ 0.05MG BASE/ML) A201533 002 Apr 21, 2016

AP FRESENIUS KABI USA EQ 0.25MG BASE/5ML (EQ 0.05MG BASE/ML) A206801 001 Sep 19, 2018

AP EQ 0.25MG BASE/5ML (EQ 0.05MG BASE/ML) A206802 001 Sep 19, 2018

AP HOSPIRA INC EQ 0.25MG BASE/5ML (EQ 0.05MG BASE/ML) A207005 001 Sep 19, 2018

AP EQ 0.075MG BASE/1.5ML (EQ 0.05MG BASE/ML) A207005 002 Sep 19, 2018

AP MYLAN INSTITUTIONAL EQ 0.25MG BASE/5ML (EQ 0.05MG BASE/ML) A206416 001 Sep 19, 2018

AP QILU EQ 0.075MG BASE/1.5ML (EQ 0.05MG BASE/ML) A205648 002 Sep 19, 2018

AP EQ 0.25MG BASE/5ML (EQ 0.05MG BASE/ML) A205648 001 Sep 19, 2018

AP SAGENT PHARMS INC EQ 0.25MG BASE/5ML (EQ 0.05MG BASE/ML) A204289 001 Sep 19, 2018

AP EQ 0.25MG BASE/5ML (EQ 0.05MG BASE/ML) A205870 001 Sep 19, 2018

AP SANDOZ INC EQ 0.25MG BASE/5ML (EQ 0.05MG BASE/ML) A202521 001 Oct 13, 2015

AP TEVA PHARMS USA EQ 0.075MG BASE/1.5ML (EQ 0.05MG BASE/ML) A090713 002 Mar 23, 2018

AP EQ 0.25MG BASE/5ML (EQ 0.05MG BASE/ML) A090713 001 Mar 23, 2018

AP VIRTUS PHARM EQ 0.25MG BASE/5ML (EQ 0.05MG BASE/ML) A209287 001 Sep 19, 2018

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

PAMIDRONATE DISODIUM

INJECTABLE; INJECTION

PAMIDRONATE DISODIUM

AP AREVA PHARMS 30MG/VIAL A077433 001 Nov 26, 2008

AP 90MG/VIAL A077433 003 Nov 26, 2008

AP DR REDDYS 30MG/10ML (3MG/ML) A078156 001 Aug 19, 2008

AP 60MG/10ML (6MG/ML) A078156 002 Aug 19, 2008

AP 90MG/10ML (9MG/ML) A078156 003 Aug 19, 2008

AP FRESENIUS KABI USA 30MG/VIAL A075773 001 May 06, 2002

AP 30MG/10ML (3MG/ML) A076207 001 May 17, 2002

AP 90MG/VIAL A075773 002 May 06, 2002

AP 90MG/10ML (9MG/ML) A076207 002 May 17, 2002

AP ! HOSPIRA 30MG/10ML (3MG/ML) A075841 001 Jun 27, 2002

AP ! 60MG/10ML (6MG/ML) A075841 002 Jun 27, 2002

AP ! 90MG/10ML (9MG/ML) A075841 003 Jun 27, 2002

AP MYLAN LABS LTD 30MG/10ML (3MG/ML) A078520 001 Oct 31, 2008

AP 90MG/10ML (9MG/ML) A078520 002 Oct 31, 2008

AP SAGENT PHARMS INC 30MG/10ML (3MG/ML) A078373 001 Dec 23, 2008

AP 90MG/10ML (9MG/ML) A078373 002 Dec 23, 2008

AP TEVA PHARMS USA 30MG/10ML (3MG/ML) A076153 001 Mar 27, 2002

AP 90MG/10ML (9MG/ML) A076153 002 Mar 27, 2002

AP WEST-WARD PHARMS INT 30MG/VIAL A075290 001 Apr 30, 2001

AP +! 30MG/10ML (3MG/ML) N021113 001 Mar 04, 2002

AP 90MG/VIAL A075290 003 Apr 30, 2001

AP +! 90MG/10ML (9MG/ML) N021113 002 Mar 04, 2002

AREVA PHARMS 60MG/VIAL A077433 002 Nov 26, 2008

PRESCRIPTION DRUG PRODUCT LIST

PANCRELIPASE (AMYLASE;LIPASE;PROTEASE)

CAPSULE, DELAYED RELEASE;ORAL

CREON

+	ABBVIE	60,000USP UNITS;12,000USP UNITS;38,000USP UNITS	N020725 002	Apr 30, 2009
+		15,000USP UNITS;3,000USP UNITS;9,500USP UNITS	N020725 004	Jul 12, 2011
+		30,000USP UNITS;6,000USP UNITS;19,000USP UNITS	N020725 001	Apr 30, 2009
+		180,000USP UNITS;36,000USP UNITS;114,000USP UNITS	N020725 005	Mar 14, 2013
+	!	120,000USP UNITS;24,000USP UNITS;76,000USP UNITS	N020725 003	Apr 30, 2009

PANCREAZE

+	VIVUS INC	10,850USP UNITS;2,600USP UNITS;6,200USP UNITS	N022523 005	Mar 07, 2014
+		24,600USP UNITS; 4,200USP UNITS; 14,200USP UNITS	N022523 001	Apr 12, 2010
+		61,500USP UNITS; 10,500USP UNITS; 35,500USP UNITS	N022523 002	Apr 12, 2010
+		83,900USP UNITS; 21,000USP UNITS; 54,700USP UNITS	N022523 004	Apr 12, 2010
+	!	98,400USP UNITS; 16,800USP UNITS; 56,800USP UNITS	N022523 003	Apr 12, 2010

PERTZYE

+	DIGESTIVE CARE INC	30,250USP UNITS;8,000USP UNITS;28,750USP UNITS	N022175 001	May 17, 2012
+	!	60,500USP UNITS;16,000USP UNITS;57,500USP UNITS	N022175 002	May 17, 2012
+		15,125USP UNITS;4,000USP UNITS;14,375USP UNITS	N022175 003	Oct 06, 2016
+		90,750USP UNITS;24,000USP UNITS;86,250USP UNITS	N022175 004	Jul 13, 2017

ZENPEP

+	ZENPEP	168,000USP UNITS;40,000USP UNITS;126,000USP UNITS	N022210 007	Mar 25, 2014
+		14,000USP UNITS;3,000USP UNITS;10,000USP UNITS	N022210 005	Jun 15, 2011
+		24,000USP UNITS;5,000USP UNITS;17,000USP UNITS	N022210 001	Aug 27, 2009
+		42,000USP UNITS;10,000USP UNITS;32,000USP UNITS	N022210 002	Aug 27, 2009
+		63,000USP UNITS;15,000USP UNITS;47,000USP UNITS	N022210 003	Aug 27, 2009
+		84,000USP UNITS;20,000USP UNITS;63,000USP UNITS	N022210 004	Aug 27, 2009
+	!	105,000USP UNITS;25,000USP UNITS;79,000USP UNITS	N022210 006	Jul 13, 2011

TABLET;ORAL

VIOKACE

+	VIOKACE	39,150USP UNITS;10,440USP UNITS;39,150USP UNITS	N022542 001	Mar 01, 2012
+	!	78,300USP UNITS;20,880USP UNITS;78,300USP UNITS	N022542 002	Mar 01, 2012

PANCURONIUM BROMIDE

INJECTABLE;INJECTION

PANCURONIUM BROMIDE

AP	!	DR REDDYS	1MG/ML	A072759 001	Jul 31, 1990
AP		HOSPIRA	1MG/ML	A072320 001	Jan 19, 1989
	!	DR REDDYS	2MG/ML	A072760 001	Jul 31, 1990

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

FOR SUSPENSION, DELAYED RELEASE;ORAL

PROTONIX

+	WYETH PHARMS	EQ 40MG BASE	N022020 001	Nov 14, 2007
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INJECTABLE;IV (INFUSION)

PANTOPRAZOLE SODIUM

AP		AKORN INC	EQ 40MG BASE/VIAL	A079197 001	Nov 08, 2012
AP		AUROBINDO PHARMA LTD	EQ 40MG BASE/VIAL	A205675 001	Mar 30, 2016
AP		SANDOZ INC	EQ 40MG BASE/VIAL	A090296 001	Jul 14, 2015

PRESCRIPTION DRUG PRODUCT LIST

PARICALCITOL

CAPSULE; ORAL

ZEMPLAR

AB + **2MCG** **N021606 002** May 26, 2005
SOLUTION; INTRAVENOUS

PARICALCITOL

AP	ACCORD HLTHCARE	<u>0.002MG/ML (0.002MG/ML)</u>	N207174 001	Feb 04, 2016
AP		<u>0.005MG/ML (0.005MG/ML)</u>	N207174 002	Feb 04, 2016
AP		<u>0.01MG/2ML (0.005MG/ML)</u>	N207174 003	Feb 04, 2016
AP	AKORN	<u>0.005MG/ML (0.005MG/ML)</u>	A207692 001	Oct 16, 2017
AP	AMNEAL PHARMS CO	<u>0.002MG/ML (0.002MG/ML)</u>	A206699 001	Mar 09, 2017
AP		<u>0.005MG/ML (0.005MG/ML)</u>	A206699 002	Mar 09, 2017
AP		<u>0.01MG/2ML (0.005MG/ML)</u>	A206699 003	Mar 09, 2017
AP	AUROBINDO PHARMA LTD	<u>0.002MG/ML (0.002MG/ML)</u>	A205982 001	Oct 09, 2018
AP		<u>0.005MG/ML (0.005MG/ML)</u>	A205982 002	Oct 09, 2018
AP		<u>0.01MG/2ML (0.005MG/ML)</u>	A205982 003	Oct 09, 2018
AP	DR REDDYS	<u>0.002MG/ML (0.002MG/ML)</u>	A204910 001	Aug 17, 2016
AP		<u>0.005MG/ML (0.005MG/ML)</u>	A204910 002	Aug 17, 2016
AP		<u>0.01MG/2ML (0.005MG/ML)</u>	A204910 003	Aug 17, 2016
AP	HIKMA PHARMS	<u>0.002MG/ML (0.002MG/ML)</u>	N205917 001	Nov 18, 2014
AP		<u>0.005MG/ML (0.005MG/ML)</u>	N205917 002	Nov 18, 2014
AP		<u>0.01MG/2ML (0.005MG/ML)</u>	N205917 003	Nov 18, 2014
AP	HOSPIRA INC	<u>0.002MG/ML (0.002MG/ML)</u>	N201657 001	Oct 21, 2014
AP		<u>0.005MG/ML (0.005MG/ML)</u>	N201657 002	Oct 21, 2014
AP		<u>0.01MG/2ML (0.005MG/ML)</u>	N201657 003	Oct 21, 2014
AP	MYLAN LABS LTD	<u>0.002MG/ML (0.002MG/ML)</u>	A203897 001	Nov 02, 2017
AP		<u>0.005MG/ML (0.005MG/ML)</u>	A203897 002	Nov 02, 2017
AP		<u>0.01MG/2ML (0.005MG/ML)</u>	A203897 003	Nov 02, 2017
AP	SANDOZ INC	<u>0.002MG/ML (0.002MG/ML)</u>	A091108 001	Jul 27, 2011
AP		<u>0.005MG/ML (0.005MG/ML)</u>	A091108 002	Jul 27, 2011
AP		<u>0.01MG/2ML (0.005MG/ML)</u>	A091108 003	Jul 27, 2011

ZEMPLAR

AP	+!	ABEVIE	<u>0.002MG/ML (0.002MG/ML)</u>	N020819 002	Feb 01, 2000
AP	+!		<u>0.005MG/ML (0.005MG/ML)</u>	N020819 001	Apr 17, 1998
AP	+!		<u>0.01MG/2ML (0.005MG/ML)</u>	N020819 003	Feb 01, 2000

PAROMOMYCIN SULFATE

CAPSULE; ORAL

PAROMOMYCIN SULFATE

AA	HERITAGE PHARMS INC	<u>EQ 250MG BASE</u>	A065173 001	Dec 14, 2007
AA	! SUN PHARM INDS INC	<u>EQ 250MG BASE</u>	A064171 001	Jun 30, 1997

PAROXETINE HYDROCHLORIDE

SUSPENSION; ORAL

PAXIL

+! APOTEX TECHNOLOGIES EQ 10MG BASE/5ML N020710 001 Jun 25, 1997

TABLET; ORAL

PAROXETINE

AB	PRINSTON INC	<u>EQ 10MG BASE</u>	A203854 001	Oct 31, 2014
AB		<u>EQ 20MG BASE</u>	A203854 002	Oct 31, 2014
AB		<u>EQ 30MG BASE</u>	A203854 003	Oct 31, 2014
AB		<u>EQ 40MG BASE</u>	A203854 004	Oct 31, 2014

PAROXETINE HYDROCHLORIDE

AB	APOTEX	<u>EQ 10MG BASE</u>	A075356 001	Jul 30, 2003
AB		<u>EQ 20MG BASE</u>	A075356 002	Jul 30, 2003
AB		<u>EQ 30MG BASE</u>	A075356 003	Jul 30, 2003
AB		<u>EQ 40MG BASE</u>	A075356 004	Jul 30, 2003
AB	AUROBINDO PHARMA	<u>EQ 10MG BASE</u>	A078406 001	Jul 25, 2007
AB		<u>EQ 20MG BASE</u>	A078406 002	Jul 25, 2007
AB		<u>EQ 30MG BASE</u>	A078406 003	Jul 25, 2007
AB		<u>EQ 40MG BASE</u>	A078406 004	Jul 25, 2007
AB	MYLAN	<u>EQ 10MG BASE</u>	A078902 001	Mar 13, 2008
AB		<u>EQ 20MG BASE</u>	A078902 002	Mar 13, 2008
AB		<u>EQ 30MG BASE</u>	A078902 003	Mar 13, 2008
AB		<u>EQ 40MG BASE</u>	A078902 004	Mar 13, 2008
AB	OXFORD PHARMS	<u>EQ 10MG BASE</u>	A076968 001	Jun 21, 2010
AB		<u>EQ 20MG BASE</u>	A076968 002	Jun 21, 2010
AB		<u>EQ 30MG BASE</u>	A076968 003	Jun 21, 2010
AB		<u>EQ 40MG BASE</u>	A076968 004	Jun 21, 2010
AB	TEVA	<u>EQ 10MG BASE</u>	A076618 001	Aug 15, 2005
AB		<u>EQ 20MG BASE</u>	A076618 002	Aug 15, 2005
AB		<u>EQ 30MG BASE</u>	A076618 003	Aug 15, 2005
AB		<u>EQ 40MG BASE</u>	A076618 004	Aug 15, 2005

PRESCRIPTION DRUG PRODUCT LIST

PAROXETINE HYDROCHLORIDE

TABLET; ORAL

PAROXETINE HYDROCHLORIDE

<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 TECHNOLOGIES	<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>	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>	MYLAN	<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>	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

PAXIL CR

<u>AB</u>	+	APOTEX TECHNOLOGIES	<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>	+	!	SEBELA IRELAND LTD	<u>EQ 7.5MG BASE</u>	<u>N204516 001</u>	Jun 28, 2013
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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

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

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

PATIROMER SORBITE X CALCIUM

POWDER; ORAL

VELTASSA

+	RELYPSA INC	EQ 8.4GM BASE/PACKET	N205739 001	Oct 21, 2015
+		EQ 16.8GM BASE/PACKET	N205739 002	Oct 21, 2015
+	!	EQ 25.2GM BASE/PACKET	N205739 003	Oct 21, 2015

PATISIRAN SODIUM

SOLUTION; INTRAVENOUS

ONPATRO

+	!	ALNYLAM PHARMS INC	EQ 10MG BASE/5ML (EQ 2MG BASE/ML)	N210922 001	Aug 10, 2018
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PRESCRIPTION DRUG PRODUCT LISTPAZOPANIB HYDROCHLORIDE

TABLET; ORAL

VOTRIENT

+! NOVARTIS EQ 200MG BASE N022465 001 Oct 19, 2009

PEGAPTANIB SODIUM

INJECTABLE; INTRAVITREAL

MACUGEN

+! VALEANT PHARMS LLC EQ 0.3MG ACID/0.09ML N021756 001 Dec 17, 2004

PEGVISOMANT

INJECTABLE; SUBCUTANEOUS

SOMAVERT

+! PHARMACIA 10MG/VIAL N021106 001 Mar 25, 2003

+! 15MG/VIAL N021106 002 Mar 25, 2003

+! 20MG/VIAL N021106 003 Mar 25, 2003

+! 25MG/VIAL N021106 004 Jul 31, 2014

+! 30MG/VIAL N021106 005 Jul 31, 2014

PEMETREXED

SOLUTION; INTRAVENOUS

PEMFEXY

+! EAGLE PHARMS 500MG/20ML (25MG/ML) N209472 001 Feb 08, 2020

PEMETREXED DISODIUM

POWDER; INTRAVENOUS

ALIMTA

+! LILLY EQ 100MG BASE/VIAL N021462 002 Sep 07, 2007

+! EQ 500MG BASE/VIAL N021462 001 Feb 04, 2004

PENCICLOVIR

CREAM; TOPICAL

DENA VIR

+! MYLAN 1% N020629 001 Sep 24, 1996

PENICILLAMINE

CAPSULE; ORAL

CUPRIMINE**AB** +! VALEANT PHARMS INTL **250MG** **N019853 001**PENICILLAMINE**AB** ANI PHARMS INC **250MG** **A209921 001** May 07, 2019**AB** PAR PHARM INC **250MG** **A211231 001** Dec 23, 2019**AB** WATSON LABS INC **250MG** **A210976 001** Jun 24, 2019

TABLET; ORAL

DEPEN**AB** +! MYLAN SPECIALITY LP **250MG** **N019854 001**PENICILLAMINE**AB** PAR PHARM INC **250MG** **A211196 001** Dec 23, 2019**AB** TEVA PHARMS USA **250MG** **A211497 001** Feb 13, 2020PENICILLIN 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** HANFORD GC **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**AP** **20,000,000 UNITS/VIAL** **A065079 003** Aug 30, 2002PFIZERPEN**AP** ! PFIZER **5,000,000 UNITS/VIAL** **A060657 002****AP** ! **20,000,000 UNITS/VIAL** **A060657 003**

PRESCRIPTION DRUG PRODUCT LIST

PENICILLIN G POTASSIUM

INJECTABLE; INJECTION

PENICILLIN G POTASSIUM

HANFORD GC 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 V POTASSIUMAA DAVA PHARMS INC EQ 125MG BASE/5ML A062981 001 Feb 10, 1989AA EQ 250MG BASE/5ML A062981 002 Feb 10, 1989PENICILLIN-VKAA TEVA EQ 125MG BASE/5ML A060456 001AA ! EQ 250MG BASE/5ML A060456 002

TABLET; ORAL

PENICILLIN V POTASSIUMAB AUROBINDO PHARMA EQ 250MG BASE A065435 001 Apr 29, 2008AB EQ 500MG BASE A065435 002 Apr 29, 2008AB DAVA PHARMS INC EQ 250MG BASE A062936 001 Nov 25, 1988AB EQ 500MG BASE A062935 001 Nov 23, 1988AB HIKMA PHARMS EQ 250MG BASE A090549 001 Oct 11, 2013AB EQ 500MG BASE A090549 002 Oct 11, 2013AB SANDOZ EQ 250MG BASE A064071 001 Nov 30, 1995AB ! EQ 500MG BASE A064071 002 Nov 30, 1995PENICILLIN-VKAB TEVA EQ 250MG BASE A060711 002AB EQ 500MG BASE A060711 003PENTAMIDINE ISETHIONATE

FOR SOLUTION; INHALATION

NEBUPENTAN +! FRESENIUS KABI USA 300MG/VIAL N019887 001 Jun 15, 1989PENTAMIDINE ISETHIONATEAN SETON PHARMS 300MG/VIAL A206667 001 Apr 24, 2019

INJECTABLE; INJECTION

PENTAMAP +! FRESENIUS KABI USA 300MG/VIAL N019264 001 Oct 16, 1984PENTAMIDINE ISETHIONATEAP SETON PHARMS 300MG/VIAL A206666 001 Sep 28, 2017PENTETATE CALCIUM TRISODIUM

SOLUTION; INHALATION, INTRAVENOUS

PENTETATE CALCIUM TRISODIUM

+! HAMELN PHARMA PLUS EQ 1GM BASE/5ML (EQ 200MG BASE/ML) N021749 001 Aug 11, 2004

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 SODIUM

INJECTABLE; INJECTION

NEMBUTAL SODIUMAP ! OAK PHARMS 50MG/ML A083246 001PENTOBARBITAL SODIUMAP ATHENEX 50MG/ML A206677 001 Nov 27, 2017AP CUSTOPHARM INC 50MG/ML A203619 001 Nov 13, 2017AP SAGENT PHARMS INC 50MG/ML A206404 001 May 23, 2016

PRESCRIPTION DRUG PRODUCT LISTPENTOSAN POLYSULFATE SODIUM

CAPSULE; ORAL

ELMIRON

+! JANSSEN PHARMS 100MG N020193 001 Sep 26, 1996

PENTOSTATIN

INJECTABLE; INJECTION

NIPENT**AP** +! HOSPIRA INC **10MG/VIAL** **N020122 001** Oct 11, 1991PENTOSTATIN**AP** WEST-WARD PHARMS **10MG/VIAL** **A077841 001** Aug 07, 2007
INTPENTOXIFYLLINE

TABLET, EXTENDED RELEASE; ORAL

PENTOXIFYLLINE**AB** ! APOTEX **400MG** **A075191 001** Jun 09, 1999**AB** VALEANT PHARMS **400MG** **A075028 001** Jul 20, 1998PENTOXIL**AB** UPSHER SMITH LABS **400MG** **A074962 001** Mar 31, 1999PERAMIVIR

SOLUTION; INTRAVENOUS

RAPIVAB

+! BIOCRYST 200MG/20ML (10MG/ML) N206426 001 Dec 19, 2014

PERAMPANEL

SUSPENSION; ORAL

FYCOMPA

+! EISAI INC 0.5MG/ML N208277 001 Apr 29, 2016

TABLET; ORAL

FYCOMPA

+ EISAI INC 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

PERFLUTREN

INJECTABLE; INTRAVENOUS

DEFINITY

+! LANTHEUS MEDCL 6.52MG/ML N021064 001 Jul 31, 2001

PERINDOPRIL ERBUMINE

TABLET; ORAL

PERINDOPRIL ERBUMINE**AB** AUROBINDO PHARMA **2MG** **A079070 001** Nov 10, 2009**AB** **4MG** **A079070 002** Nov 10, 2009**AB** ! **8MG** **A079070 003** Nov 10, 2009**AB** HIKMA **2MG** **A090072 001** Nov 10, 2009**AB** **4MG** **A090072 002** Nov 10, 2009**AB** **8MG** **A090072 003** Nov 10, 2009PERMETHRIN

CREAM; TOPICAL

PERMETHRIN**AB** ACTAVIS LABS **5%** **A074806 001** Jan 23, 1998**AB** ENCUBE ETHICALS **5%** **A211303 001** Apr 03, 2019**AB** ! PERRIGO ISRAEL **5%** **A076369 001** Apr 21, 2003PERPHENAZINE

TABLET; ORAL

PERPHENAZINE**AB** MYLAN **2MG** **A206691 001** Apr 14, 2017**AB** **4MG** **A206691 002** Apr 14, 2017**AB** **8MG** **A206691 003** Apr 14, 2017**AB** **16MG** **A206691 004** Apr 14, 2017**AB** RISING **2MG** **A205056 001** Mar 01, 2019**AB** **4MG** **A205056 002** Mar 01, 2019**AB** **8MG** **A205056 003** Mar 01, 2019**AB** **16MG** **A205056 004** Mar 01, 2019**AB** SANDOZ **2MG** **A089685 002** Dec 08, 1988**AB** **4MG** **A089685 003** Dec 08, 1988**AB** **8MG** **A089685 001** Dec 08, 1988**AB** ! **16MG** **A089685 004** Dec 08, 1988**AB** VINTAGE PHARMS **2MG** **A040226 001** Dec 31, 1998

PRESCRIPTION DRUG PRODUCT LIST

3-348 (of 453)

PERPHENAZINE

TABLET; ORAL

PERPHENAZINE

<u>AB</u>		<u>4MG</u>	<u>A040226</u>	<u>002</u>	Dec 31, 1998
<u>AB</u>		<u>8MG</u>	<u>A040226</u>	<u>003</u>	Dec 31, 1998
<u>AB</u>		<u>16MG</u>	<u>A040226</u>	<u>004</u>	Dec 31, 1998
<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

PEXIDARTINIB HYDROCHLORIDE

CAPSULE; ORAL

TURALIO

+! DAIICHI SANKYO INC EQ 200MG BASE N211810 001 Aug 02, 2019

PHENDIMETRAZINE TARTRATE

CAPSULE, EXTENDED RELEASE; ORAL

PHENDIMETRAZINE TARTRATE

+! VIRTUS PHARMS 105MG N018074 001

TABLET; ORAL

BONTRIL PDMAA ! VALEANT 35MG A085272 001PHENDIMETRAZINE TARTRATE

<u>AA</u>	ELITE LABS INC	<u>35MG</u>	<u>A040762</u>	<u>001</u>	Jan 28, 2008
<u>AA</u>		<u>35MG</u>	<u>A203600</u>	<u>001</u>	Dec 27, 2017
<u>AA</u>	KVK TECH	<u>35MG</u>	<u>A091042</u>	<u>001</u>	Aug 31, 2010
<u>AA</u>	MIKART	<u>35MG</u>	<u>A089452</u>	<u>001</u>	Oct 30, 1991
<u>AA</u>	VIRTUS PHARMS	<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, 2010PHENOXYBENZAMINE HYDROCHLORIDE

CAPSULE; ORAL

PHENOXYBENZAMINE HYDROCHLORIDE

<u>AB</u>	HIKMA	<u>10MG</u>	<u>A201050</u>	<u>001</u>	Jul 16, 2012
<u>AB</u>	! PAR PHARM INC	<u>10MG</u>	<u>A204522</u>	<u>001</u>	Jan 24, 2017

PHENTERMINE HYDROCHLORIDE

CAPSULE; ORAL

ADIPEX-PAA ! TEVA 37.5MG A088023 001 Aug 02, 1983PHENTERMINE 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>	BARR	<u>15MG</u>	<u>A090591</u>	<u>001</u>	Mar 18, 2010
<u>AA</u>		<u>30MG</u>	<u>A090591</u>	<u>002</u>	Mar 18, 2010
<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>	ELITE LABS INC	<u>37.5MG</u>	<u>A040228</u>	<u>001</u>	Jun 19, 1997
<u>AA</u>	INVAGEN PHARMS	<u>15MG</u>	<u>A202858</u>	<u>001</u>	Feb 14, 2014
<u>AA</u>		<u>30MG</u>	<u>A202858</u>	<u>002</u>	Feb 14, 2014
<u>AA</u>		<u>30MG</u>	<u>A204414</u>	<u>001</u>	May 05, 2014
<u>AA</u>		<u>37.5MG</u>	<u>A202846</u>	<u>001</u>	Feb 05, 2014
<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
<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>	LANNETT	<u>15MG</u>	<u>A087022</u>	<u>002</u>	Jan 20, 2012
<u>AA</u>		<u>30MG</u>	<u>A087022</u>	<u>001</u>	Feb 03, 1983
<u>AA</u>	LANNETT CO INC	<u>37.5MG</u>	<u>A201961</u>	<u>001</u>	Jul 20, 2011
<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>	! SANDOZ	<u>15MG</u>	<u>A087190</u>	<u>002</u>	
<u>AA</u>	!	<u>30MG</u>	<u>A086945</u>	<u>001</u>	Jul 20, 1983

PRESCRIPTION DRUG PRODUCT LIST

PHENTERMINE HYDROCHLORIDE

CAPSULE; ORAL

PHENTERMINE HYDROCHLORIDE

AA	!		30MG	A087190	001	
AA		SUN PHARM INDUSTRIES	30MG	A040525	001	Oct 23, 2003

TABLET; ORAL

ADIPEX-P

AA	!	TEVA	37.5MG	A085128	001	
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LOMAIRA

AA	!	AVANTHI INC	8MG	A203495	001	Sep 12, 2016
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PHENTERMINE HYDROCHLORIDE

AA		AUROLIFE PHARMA LLC	37.5MG	A203068	001	Aug 06, 2014
AA		BARR	37.5MG	A090470	001	Aug 31, 2009
AA		ELITE LABS	37.5MG	A200272	001	Jan 31, 2011
AA		ELITE LABS INC	37.5MG	A040190	001	May 30, 1997
AA		INVAGEN PHARMS	37.5MG	A202942	001	Feb 05, 2014
AA		KVK TECH	37.5MG	A040876	001	Mar 31, 2008
AA		KVK TECH INC	8MG	A203436	001	Mar 17, 2017
AA		LANNETT	37.5MG	A040555	001	Apr 15, 2005
AA		NUVO PHARM	37.5MG	A205008	001	Sep 25, 2014
AA		POLYGEN PHARMS	37.5MG	A206342	001	Nov 18, 2016
AA		PRINSTON INC	37.5MG	A040377	001	Jan 04, 2002
AA		SUN PHARM INDUSTRIES	37.5MG	A040526	001	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

QSYMIA

	+	VIVUS	EQ 3.75MG BASE; 23MG	N022580	001	Jul 17, 2012
	+		EQ 7.5MG BASE; 46MG	N022580	002	Jul 17, 2012
	+		EQ 11.25MG BASE; 69MG	N022580	003	Jul 17, 2012
	+	!	EQ 15MG BASE; 92MG	N022580	004	Jul 17, 2012

PHENTOLAMINE MESYLATE

INJECTABLE; INJECTION

PHENTOLAMINE MESYLATE

AP		PRECISION DOSE INC	5MG/VIAL	A207686	001	Jul 14, 2017	
AP	!	WEST-WARD PHARMS INT	5MG/VIAL	A040235	001	Mar 11, 1998	
		ORAVERSE					
	+	!	SEPTODONT HOLDING	0.4MG/1.7ML	N022159	001	May 09, 2008

PHENYLEPHRINE HYDROCHLORIDE

SOLUTION; INTRAVENOUS

PHENYLEPHRINE HYDROCHLORIDE

AP1		AMNEAL	10MG/ML (10MG/ML)	A211079	001	Jul 05, 2018	
AP1			50MG/5ML (10MG/ML)	A211078	001	Jul 19, 2018	
AP1			100MG/10ML (10MG/ML)	A211078	002	Jul 19, 2018	
AP1		FRESENIUS KABI USA	50MG/5ML (10MG/ML)	A210666	001	Jan 30, 2019	
AP1			100MG/10ML (10MG/ML)	A210666	002	Jan 30, 2019	
AP1		MEITHEAL	50MG/5ML (10MG/ML)	A210333	001	Apr 27, 2018	
AP1			100MG/10ML (10MG/ML)	A210333	002	Apr 27, 2018	
AP1		PAR STERILE PRODUCTS	10MG/ML (10MG/ML)	A210025	001	Dec 21, 2018	
AP1			50MG/5ML (10MG/ML)	A210025	002	Dec 21, 2018	
AP1			100MG/10ML (10MG/ML)	A210025	003	Dec 21, 2018	
AP1		RICONPHARMA LLC	10MG/ML (10MG/ML)	A209967	001	Jan 16, 2020	
AP1			50MG/5ML (10MG/ML)	A209967	002	Jan 16, 2020	
AP1			100MG/10ML (10MG/ML)	A209967	003	Jan 16, 2020	
AP1		SANDOZ INC	10MG/ML (10MG/ML)	A208905	001	Jan 31, 2019	
AP1			50MG/5ML (10MG/ML)	A208905	002	Jan 31, 2019	
AP1			100MG/10ML (10MG/ML)	A208905	003	Jan 31, 2019	
		<u>VAZCULEP</u>					
AP1	+	!	AVADEL LEGACY	10MG/ML (10MG/ML)	N204300	001	Jun 27, 2014
AP1	+	!		50MG/5ML (10MG/ML)	N204300	002	Jun 27, 2014
AP1	+	!		100MG/10ML (10MG/ML)	N204300	003	Jun 27, 2014

PHENYLEPHRINE HYDROCHLORIDE

AP2		FRESENIUS KABI USA	10MG/ML (10MG/ML)	A210665	001	Jan 29, 2019
AP2		MEITHEAL	10MG/ML (10MG/ML)	A210334	001	Apr 27, 2018

PRESCRIPTION DRUG PRODUCT LIST

PHENYLEPHRINE HYDROCHLORIDE

SOLUTION; INTRAVENOUS

PHENYLEPHRINE HYDROCHLORIDE

AP2	+ !	WEST WARD PHARM CORP	10MG/ML (10MG/ML)	N203826 001	Dec 20, 2012
AP2	+ !		50MG/5ML (10MG/ML)	N203826 002	Jun 19, 2019
AP2	+ !		100MG/10ML (10MG/ML)	N203826 003	Jun 19, 2019
		BIORPHEN			
	+ !	ETON PHARMS	0.1MG/ML (0.1MG/ML)	N212909 001	Oct 21, 2019

SOLUTION/DROPS; OPHTHALMIC

PHENYLEPHRINE HYDROCHLORIDE

AT	+ !	AKORN INC	2.5%	N207926 001	Jan 15, 2015
AT	+ !		10%	N207926 002	Jan 15, 2015
AT	+ !	PARAGON BIOTECK	2.5%	N203510 001	Mar 21, 2013
AT	+ !		10%	N203510 002	Mar 21, 2013

PHENYLEPHRINE HYDROCHLORIDE; PROMETHAZINE HYDROCHLORIDE

SYRUP; ORAL

PHENYLEPHRINE HYDROCHLORIDE AND PROMETHAZINE HYDROCHLORIDE

AA		HI TECH	5MG/5ML; 6.25MG/5ML	A040675 001	Dec 23, 2014
AA	!	VINTAGE	5MG/5ML; 6.25MG/5ML	A040654 001	Dec 07, 2006
		<u>PROMETHAZINE HYDROCHLORIDE AND PHENYLEPHRINE HYDROCHLORIDE</u>			
AA		AMNEAL PHARMS	5MG/5ML; 6.25MG/5ML	A040902 001	Aug 25, 2009

PHENYTOIN

SUSPENSION; ORAL

DILANTIN-125

AB	+ !	PARKE DAVIS	125MG/5ML	N008762 001	
		<u>PHENYTOIN</u>			
AB		TARO	125MG/5ML	A040521 001	Mar 08, 2004
AB		VISTAPHARM	125MG/5ML	A040342 001	Jan 31, 2001
AB			125MG/5ML	A040610 001	Aug 18, 2005
AB		WOCKHARDT BIO AG	125MG/5ML	A040420 001	Apr 19, 2002

TABLET, CHEWABLE; ORAL

DILANTIN

AB	!	PFIZER	50MG	A084427 001	
		<u>PHENYTOIN</u>			
AB		EPIC PHARMA LLC	50MG	A040884 001	Nov 28, 2014
AB		MYLAN PHARMS INC	50MG	A200691 001	Dec 26, 2012
AB		TARO	50MG	A200565 001	Apr 17, 2014

PHENYTOIN SODIUM

CAPSULE; ORAL

DILANTIN

AB	!	PARKE-DAVIS	100MG EXTENDED	A084349 002	
		<u>EXTENDED PHENYTOIN SODIUM</u>			
AB		AMNEAL PHARMS NY	100MG EXTENDED	A040765 001	Nov 12, 2008
AB		LUPIN LTD	100MG EXTENDED	A211633 001	Sep 30, 2019
AB		MYLAN	100MG EXTENDED	A040298 001	Dec 28, 1998
AB		SUN PHARM INDS	200MG EXTENDED	A040731 001	Jun 30, 2008
AB			300MG EXTENDED	A040731 002	Jun 30, 2008
AB		TARO	100MG EXTENDED	A040684 001	Sep 05, 2006
		<u>PHENYTEK</u>			
AB		MYLAN	200MG EXTENDED	A040298 002	Dec 06, 2001
AB	!		300MG EXTENDED	A040298 003	Dec 06, 2001

PHENYTOIN SODIUM

AB		AUROBINDO PHARMA LTD	100MG EXTENDED	A204309 001	Jun 10, 2015
		<u>DILANTIN</u>			
	!	PARKE-DAVIS	30MG EXTENDED	A084349 001	

INJECTABLE; INJECTION

PHENYTOIN SODIUM

AP		ACELLA	50MG/ML	A040573 001	Sep 13, 2006
AP	!	WEST-WARD PHARMS INT	50MG/ML	A084307 001	

PHYTONADIONE

INJECTABLE; INJECTION

PHYTONADIONE

AB		DR REDDYS LABS LTD	10MG/ML	A207719 001	May 22, 2019
		<u>VITAMIN K1</u>			
AB	!	HOSPIRA	10MG/ML	A087955 001	Jul 25, 1983
		<u>PHYTONADIONE</u>			
BP	!	INTL MEDICATION	1MG/0.5ML	A083722 001	

PRESCRIPTION DRUG PRODUCT LIST

PHYTONADIONEINJECTABLE; INJECTION
VITAMIN K1BP ! HOSPIRA 1MG/0.5ML A087954 001 Jul 25, 1983
TABLET; ORALMEPHYTON**AB** +! BAUSCH **5MG** **N010104 003**PHYTONADIONE**AB** AMNEAL PHARMS CO **5MG** **A209373 001** May 11, 2018
AB ZYDUS **5MG** **A210189 001** Feb 20, 2019PILOCARPINE HYDROCHLORIDE

SOLUTION; OPHTHALMIC

ISOPTO CARPINE**AT** + NOVARTIS **1%** **N200890 001** Jun 22, 2010
AT + **2%** **N200890 002** Jun 22, 2010
AT +! **4%** **N200890 003** Jun 22, 2010PILOCARPINE HYDROCHLORIDE**AT** AKORN INC **1%** **A204398 001** Sep 27, 2017
AT **2%** **A204398 002** Sep 27, 2017
AT **4%** **A204398 003** Sep 27, 2017
AT SOMERSET THERAPS **1%** **A210384 001** Nov 25, 2019
LLC
AT **2%** **A210384 002** Nov 25, 2019
AT **4%** **A210384 003** Nov 25, 2019

TABLET; ORAL

PILOCARPINE HYDROCHLORIDE**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 PERRIGO PHARMA INTL **5MG** **A076746 001** Nov 16, 2004SALAGEN**AB** + CONCORDIA **5MG** **N020237 001** Mar 22, 1994
AB +! **7.5MG** **N020237 002** Apr 18, 2003PIMAVANSERIN 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, 2001PIMECROLIMUS**AB** ACTAVIS LABS UT INC **1%** **A209345 001** Dec 27, 2018
AB GLENMARK PHARMS LTD **1%** **A211769 001** Aug 29, 2019PIMOZIDE

TABLET; ORAL

PIMOZIDE

PAR PHARM 1MG A204521 001 Sep 28, 2015

! 2MG A204521 002 Sep 28, 2015

PINDOLOL

TABLET; ORAL

PINDOLOL**AB** ANI PHARMS INC **5MG** **A073609 002** Mar 29, 1993
AB **10MG** **A073609 001** Mar 29, 1993
AB BAYSHORE PHARMS LLC **5MG** **A211712 001** Aug 01, 2019
AB **10MG** **A211712 002** Aug 01, 2019
AB MYLAN PHARMS INC **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 **5MG** **A074063 001** Jan 27, 1994
INDUSTRIES
AB **10MG** **A074063 002** Jan 27, 1994

PRESCRIPTION DRUG PRODUCT LIST

PIOGLITAZONE HYDROCHLORIDE

TABLET; ORAL

ACTOS

AB	+	TAKEDA PHARMS USA	<u>EQ 15MG BASE</u>	<u>N021073 001</u>	Jul 15, 1999
AB	+		<u>EQ 30MG BASE</u>	<u>N021073 002</u>	Jul 15, 1999
AB	+		<u>EQ 45MG BASE</u>	<u>N021073 003</u>	Jul 15, 1999

PIOGLITAZONE HYDROCHLORIDE

AB		ACCORD HLTHCARE	<u>EQ 15MG BASE</u>	<u>A200044 001</u>	Feb 13, 2013
AB			<u>EQ 30MG BASE</u>	<u>A200044 002</u>	Feb 13, 2013
AB			<u>EQ 45MG BASE</u>	<u>A200044 003</u>	Feb 13, 2013
AB		AUROBINDO PHARMA LTD	<u>EQ 15MG BASE</u>	<u>A200268 001</u>	Feb 13, 2013
AB			<u>EQ 30MG BASE</u>	<u>A200268 002</u>	Feb 13, 2013
AB			<u>EQ 45MG BASE</u>	<u>A200268 003</u>	Feb 13, 2013
AB		BRECKENRIDGE	<u>EQ 15MG BASE</u>	<u>A078472 001</u>	Feb 13, 2013
AB			<u>EQ 30MG BASE</u>	<u>A078472 002</u>	Feb 13, 2013
AB			<u>EQ 45MG BASE</u>	<u>A078472 003</u>	Feb 13, 2013
AB		CELLTRION	<u>EQ 15MG BASE</u>	<u>A076798 001</u>	Oct 26, 2012
AB			<u>EQ 30MG BASE</u>	<u>A076798 002</u>	Oct 26, 2012
AB			<u>EQ 45MG BASE</u>	<u>A076798 003</u>	Oct 26, 2012
AB		LUPIN LTD	<u>EQ 15MG BASE</u>	<u>A204133 001</u>	Apr 07, 2014
AB			<u>EQ 30MG BASE</u>	<u>A204133 002</u>	Apr 07, 2014
AB			<u>EQ 45MG BASE</u>	<u>A204133 003</u>	Apr 07, 2014
AB		MACLEODS PHARMS LTD	<u>EQ 15MG BASE</u>	<u>A202467 001</u>	Feb 06, 2013
AB			<u>EQ 30MG BASE</u>	<u>A202467 002</u>	Feb 06, 2013
AB			<u>EQ 45MG BASE</u>	<u>A202467 003</u>	Feb 06, 2013
AB		NEOPHARMA	<u>EQ 15MG BASE</u>	<u>A078383 001</u>	Mar 12, 2013
AB			<u>EQ 30MG BASE</u>	<u>A078383 002</u>	Mar 12, 2013
AB			<u>EQ 45MG BASE</u>	<u>A078383 003</u>	Mar 12, 2013
AB		PRINSTON INC	<u>EQ 15MG BASE</u>	<u>A207806 001</u>	Apr 17, 2018
AB			<u>EQ 30MG BASE</u>	<u>A207806 002</u>	Apr 17, 2018
AB			<u>EQ 45MG BASE</u>	<u>A207806 003</u>	Apr 17, 2018
AB		PURACAP PHARM LLC	<u>EQ 15MG BASE</u>	<u>A206738 001</u>	Oct 06, 2017
AB			<u>EQ 30MG BASE</u>	<u>A206738 002</u>	Oct 06, 2017
AB			<u>EQ 45MG BASE</u>	<u>A206738 003</u>	Oct 06, 2017
AB		SANDOZ	<u>EQ 15MG BASE</u>	<u>A078670 001</u>	Feb 13, 2013
AB			<u>EQ 30MG BASE</u>	<u>A078670 002</u>	Feb 13, 2013
AB			<u>EQ 45MG BASE</u>	<u>A078670 003</u>	Feb 13, 2013
AB		TEVA PHARMS USA	<u>EQ 15MG BASE</u>	<u>A077210 001</u>	Jan 10, 2014
AB			<u>EQ 30MG BASE</u>	<u>A077210 002</u>	Jan 10, 2014
AB			<u>EQ 45MG BASE</u>	<u>A077210 003</u>	Jan 10, 2014
AB		TORRENT PHARMS LTD	<u>EQ 15MG BASE</u>	<u>A091298 001</u>	Feb 13, 2013
AB			<u>EQ 30MG BASE</u>	<u>A091298 002</u>	Feb 13, 2013
AB			<u>EQ 45MG BASE</u>	<u>A091298 003</u>	Feb 13, 2013
AB		ZYDUS PHARMS USA INC	<u>EQ 15MG BASE</u>	<u>A202456 001</u>	Feb 13, 2013
AB			<u>EQ 30MG BASE</u>	<u>A202456 002</u>	Feb 13, 2013
AB			<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

AP	APOLLO	<u>EQ 2GM BASE/VIAL;EQ 250MG BASE/VIAL</u>	<u>A207847 001</u>	Jan 13, 2017
AP		<u>EQ 3GM BASE/VIAL;EQ 375MG BASE/VIAL</u>	<u>A207847 002</u>	Jan 13, 2017
AP		<u>EQ 36GM BASE/VIAL;EQ 4.5GM BASE/VIAL</u>	<u>A207848 002</u>	Jan 13, 2017
AP		<u>EQ 4GM BASE/VIAL;EQ 500MG BASE/VIAL</u>	<u>A207847 003</u>	Jan 13, 2017
AP		<u>EQ 12GM BASE/VIAL;EQ 1.5GM BASE/VIAL</u>	<u>A207848 001</u>	May 11, 2018
AP	AUROBINDO PHARMA LTD	<u>EQ 2GM BASE/VIAL;EQ 250MG BASE/VIAL</u>	<u>A065498 001</u>	May 23, 2011
AP		<u>EQ 3GM BASE/VIAL;EQ 375MG BASE/VIAL</u>	<u>A065498 002</u>	May 23, 2011
AP		<u>EQ 4GM BASE/VIAL;EQ 500MG BASE/VIAL</u>	<u>A065498 003</u>	May 23, 2011
AP	FRESENIUS KABI	<u>EQ 2GM BASE/VIAL;EQ 250MG BASE/VIAL</u>	<u>A203719 001</u>	May 18, 2018
AP		<u>EQ 3GM BASE/VIAL;EQ 375MG BASE/VIAL</u>	<u>A203719 002</u>	May 18, 2018
AP		<u>EQ 4GM BASE/VIAL;EQ 500MG BASE/VIAL</u>	<u>A203719 003</u>	May 18, 2018
AP		<u>EQ 36GM BASE/VIAL;EQ 4.5GM BASE/VIAL</u>	<u>A203720 001</u>	May 11, 2018

PRESCRIPTION DRUG PRODUCT LIST

PIPERACILLIN SODIUM; TAZOBACTAM SODIUM

INJECTABLE; INJECTION

PIPERACILLIN AND TAZOBACTAM

<u>AP</u>	FRESENIUS KABI USA	<u>EQ 12GM BASE/VIAL;EQ 1.5GM BASE/VIAL</u>	<u>A206204 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>	MYLAN LABS LTD	<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
<u>AP</u>		<u>EQ 4GM BASE/VIAL;EQ 500MG BASE/VIAL</u>	<u>A065458 003</u>	Aug 15, 2014
<u>AP</u>	QILU	<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>	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>	SANDOZ INC	<u>EQ 12GM BASE/VIAL;EQ 1.5GM BASE/VIAL</u>	<u>A203557 001</u>	Oct 29, 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
<u>ZOSYN</u>				
<u>AP</u>	+! WYETH PHARMS	<u>EQ 2GM BASE/VIAL;EQ 250MG BASE/VIAL</u>	<u>N050684 001</u>	Oct 22, 1993
<u>AP</u>	+!	<u>EQ 3GM BASE/VIAL;EQ 375MG BASE/VIAL</u>	<u>N050684 002</u>	Oct 22, 1993
<u>AP</u>	+!	<u>EQ 4GM BASE/VIAL;EQ 500MG BASE/VIAL</u>	<u>N050684 003</u>	Oct 22, 1993
<u>AP</u>	+!	<u>EQ 36GM BASE/VIAL;EQ 4.5GM BASE/VIAL</u>	<u>N050684 004</u>	Oct 22, 1993

ZOSYN IN PLASTIC CONTAINER

+!	WYETH PHARMS	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

+!	GENENTECH INC	267MG	N022535 001	Oct 15, 2014
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TABLET; ORAL

ESBRIET

+	GENENTECH INC	267MG	N208780 001	Jan 11, 2017
+!		801MG	N208780 003	Jan 11, 2017

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>	CADILA	<u>10MG</u>	<u>A205585 001</u>	Jul 17, 2018
<u>AB</u>		<u>20MG</u>	<u>A205585 002</u>	Jul 17, 2018
<u>AB</u>	FLAMINGO PHARMS	<u>10MG</u>	<u>A207938 001</u>	Sep 09, 2016
<u>AB</u>		<u>20MG</u>	<u>A207938 002</u>	Sep 09, 2016
<u>AB</u>	HIKMA PHARMS	<u>10MG</u>	<u>A209256 001</u>	Aug 11, 2017
<u>AB</u>		<u>20MG</u>	<u>A209256 002</u>	Aug 11, 2017
<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>A074116 001</u>	Jun 15, 1993
<u>AB</u>		<u>20MG</u>	<u>A074118 001</u>	Jun 15, 1993
<u>AB</u>	PII	<u>10MG</u>	<u>A206136 001</u>	Jun 20, 2017
<u>AB</u>		<u>20MG</u>	<u>A206136 002</u>	Jun 20, 2017
<u>AB</u>	STRIDES PHARMA	<u>10MG</u>	<u>A210347 001</u>	Jan 26, 2018
<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 LABS LTD	<u>10MG</u>	<u>A208340 001</u>	Apr 13, 2017
<u>AB</u>		<u>20MG</u>	<u>A208340 002</u>	Apr 13, 2017

PRESCRIPTION DRUG PRODUCT LIST

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		AUROBINDO PHARMA LTD	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		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

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

PLAZOMICIN SULFATE

SOLUTION;INTRAVENOUS

ZEMDRI

	+	CIPLA USA	EQ 500MG BASE/10ML (EQ 50MG BASE/ML)	N210303 001	Jun 25, 2018
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PLECANATIDE

TABLET;ORAL

TRULANCE

	+	SALIX	3MG	N208745 001	Jan 19, 2017
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PLERIXAFOR

SOLUTION;SUBCUTANEOUS

MOZOBIL

	+	GENZYME	24MG/1.2ML (20MG/ML)	N022311 001	Dec 15, 2008
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PODOFILOX

GEL;TOPICAL

CONDYLOX

	+	ALLERGAN	0.5%	N020529 001	Mar 13, 1997
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SOLUTION;TOPICAL

CONDYLOX

AT	+	TEVA BRANDED PHARM	0.5%	N019795 001	Dec 13, 1990
AT		PADDOCK LLC	0.5%	A075600 001	Jan 29, 2002

POLIDOCANOL

SOLUTION;INTRAVENOUS

ASCLERA

	+	CHEMISCH FBRK KRSSLR	10MG/2ML (5MG/ML)	N021201 001	Mar 30, 2010
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	+	!	20MG/2ML (10MG/ML)	N021201 002	Mar 30, 2010
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VARITHENA

	+	PROVENSIS	77.5MG/7.75ML (10MG/ML)	N205098 002	Dec 21, 2017
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	+	!	180MG/18ML (10MG/ML)	N205098 001	Nov 25, 2013
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POLYETHYLENE GLYCOL 3350; POTASSIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE

FOR SOLUTION;ORAL

LAX-LYTE WITH FLAVOR PACKS

AA		PADDOCK LLC	420GM/BOT;1.48GM/BOT;5.72GM/BOT;11.2GM/BOT	A079232 001	Feb 25, 2010
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NULYTELY

AA	+	BRAINTREE	420GM/BOT;1.48GM/BOT;5.72GM/BOT;11.2GM/BOT	N019797 001	Apr 22, 1991
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NULYTELY-FLAVORED

AA	+	BRAINTREE	420GM/BOT;1.48GM/BOT;5.72GM/BOT;11.2GM/BOT	N019797 002	Nov 18, 1994
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PRESCRIPTION DRUG PRODUCT LIST

POLYETHYLENE GLYCOL 3350; POTASSIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE

FOR SOLUTION;ORAL

PEG-3350, POTASSIUM CHLORIDE, SODIUM BICARBONATE, SODIUM CHLORIDE

AA	BRECKENRIDGE	<u>420GM/BOT;1.48GM/BOT;5.72GM/BOT;11.2GM/BOT</u>	<u>A202060 001</u>	Mar 08, 2017
AA	NOVEL LABS INC	<u>420GM/BOT;1.48GM/BOT;5.72GM/BOT;11.2GM/BOT</u>	<u>A090019 001</u>	May 27, 2009
AA	STRIDES PHARMA	<u>420GM/BOT;1.48GM/BOT;5.72GM/BOT;11.2GM/BOT</u>	<u>A204559 001</u>	Apr 13, 2015
TRILYTE				
AA	MYLAN	<u>420GM/BOT;1.48GM/BOT;5.72GM/BOT;11.2GM/BOT</u>	<u>A076491 001</u>	Feb 05, 2004

POLYETHYLENE GLYCOL 3350; POTASSIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE; SODIUM SULFATE ANHYDROUS

FOR SOLUTION;ORAL

COLYTE WITH FLAVOR PACKS

AA	MYLAN SPECIALITY LP	<u>240GM/BOT;2.98GM/BOT;6.72GM/BOT;5.84GM/BOT;22.72GM/BOT</u>	<u>N018983 012</u>	Oct 08, 1998
GOLYTELY				
AA	+! BRAINTREE	<u>236GM/BOT;2.97GM/BOT;6.74GM/BOT;5.86GM/BOT;22.74GM/BOT</u>	<u>N019011 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 001</u>	Jun 01, 2009
AA		<u>240GM/BOT;2.98GM/BOT;6.72GM/BOT;5.84GM/BOT;22.72GM/BOT</u>	<u>A090186 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 001</u>	Dec 21, 2018

POLYMYXIN B SULFATE

INJECTABLE; INJECTION

POLYMYXIN B SULFATE

AP	AUROBINDO PHARMA LTD	<u>EQ 500,000 UNITS BASE/VIAL</u>	<u>A206589 001</u>	Apr 04, 2016
AP	FRESENIUS KABI USA	<u>EQ 500,000 UNITS BASE/VIAL</u>	<u>A065372 001</u>	Jan 10, 2008
AP	! GLAND PHARMA LTD	<u>EQ 500,000 UNITS BASE/VIAL</u>	<u>A207322 001</u>	Apr 14, 2016
AP	MYLAN ASI	<u>EQ 500,000 UNITS BASE/VIAL</u>	<u>A090110 001</u>	Jun 29, 2011
AP	XELLIA PHARMS APS	<u>EQ 500,000 UNITS BASE/VIAL</u>	<u>A202766 001</u>	Jan 15, 2014
AP	XGEN PHARMS	<u>EQ 500,000 UNITS BASE/VIAL</u>	<u>A063000 001</u>	Sep 30, 1994

POLYMYXIN B SULFATE; TRIMETHOPRIM SULFATE

SOLUTION/DROPS;OPHTHALMIC

POLYTRIM

AT	+! ALLERGAN	<u>10,000 UNITS/ML;EQ 1MG BASE/ML</u>	<u>N050567 001</u>	Oct 20, 1988
TRIMETHOPRIM SULFATE AND POLYMYXIN B SULFATE				
AT	AKORN INC	<u>10,000 UNITS/ML;EQ 1MG BASE/ML</u>	<u>A065006 001</u>	Dec 17, 1998
AT	BAUSCH AND LOMB	<u>10,000 UNITS/ML;EQ 1MG BASE/ML</u>	<u>A064120 001</u>	Feb 14, 1997
AT	SANDOZ INC	<u>10,000 UNITS/ML;EQ 1MG BASE/ML</u>	<u>A064211 001</u>	Apr 13, 1998

POMALIDOMIDE

CAPSULE;ORAL

POMALYST

+	CELGENE	1MG	N204026 001	Feb 08, 2013
+		2MG	N204026 002	Feb 08, 2013
+		3MG	N204026 003	Feb 08, 2013
+	!	4MG	N204026 004	Feb 08, 2013

PONATINIB HYDROCHLORIDE

TABLET;ORAL

ICLUSIG

+	ARIAD	EQ 15MG BASE	N203469 001	Dec 14, 2012
+		EQ 30MG BASE	N203469 003	Apr 23, 2015
+	!	EQ 45MG BASE	N203469 002	Dec 14, 2012

PORACTANT ALFA

SUSPENSION; INTRATRACHEAL

CUROSURF

+	CHIESI USA INC	80MG/ML	N020744 001	Nov 18, 1999
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PORFIMER SODIUM

INJECTABLE; INJECTION

PHOTOFRIN

	PINNACLE BIOLGS	75MG/VIAL	N020451 001	Dec 27, 1995
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PRESCRIPTION DRUG PRODUCT LIST

POSACONAZOLE

SOLUTION; INTRAVENOUS

NOXAFIL

+! MERCK SHARP DOHME 300MG/16.7ML (18MG/ML) N205596 001 Mar 13, 2014

SUSPENSION; ORAL

NOXAFIL

+! SCHERING 40MG/ML N022003 001 Sep 15, 2006

TABLET, DELAYED RELEASE; ORAL

NOXAFIL**AB** +! MERCK SHARP DOHME **100MG** **N205053 001** Nov 25, 2013POSACONAZOLE**AB** SINOTHERAPEUTICS **100MG** **A212411 001** Aug 21, 2019
INCPOTASSIUM ACETATE

INJECTABLE; INJECTION

POTASSIUM ACETATE**AP** EXELA PHARMA SCS **2MEQ/ML** **A206203 001** Dec 29, 2015
LLC**AP** +! HOSPIRA **2MEQ/ML** **N018896 001** Jul 20, 1984POTASSIUM CHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

KLOR-CON**AB** UPSHER SMITH LABS **8MEQ** **A203106 001** Jul 10, 2015**AB** **10MEQ** **A203106 002** Jul 10, 2015MICRO-K**AB** + NESHER PHARMS **8MEQ** **N018238 001**MICRO-K 10**AB** + NESHER PHARMS **10MEQ** **N018238 002** May 14, 1984POTASSIUM CHLORIDE**AB** ACTAVIS LABS FL INC **8MEQ** **A077419 001** Jun 02, 2008**AB** ! **10MEQ** **A077419 002** Jun 02, 2008**AB** ADARE PHARMS INC **8MEQ** **A208864 001** Mar 17, 2017**AB** **10MEQ** **A208864 002** Mar 17, 2017**AB** AMNEAL PHARMS **10MEQ** **A202128 001** Feb 22, 2013**AB** ANCHEN PHARMS **8MEQ** **A202886 001** Dec 26, 2013**AB** **10MEQ** **A202886 002** Dec 26, 2013**AB** GLENMARK PHARMS LTD **10MEQ** **A202868 001** Jan 19, 2016**AB** LANNETT CO INC **8MEQ** **A204210 001** Mar 28, 2016**AB** **10MEQ** **A204210 002** Mar 28, 2016**AB** LUPIN LTD **8MEQ** **A203002 001** Dec 18, 2015**AB** **10MEQ** **A203002 002** Dec 18, 2015**AB** NOVEL LABS INC **8MEQ** **A204828 001** Aug 16, 2016**AB** **10MEQ** **A204828 002** Aug 16, 2016**AB** PADDOCK LLC **8MEQ** **A200185 001** May 18, 2011**AB** **10MEQ** **A200185 002** May 18, 2011**AB** PII **8MEQ** **A205549 001** Dec 08, 2015**AB** **10MEQ** **A205549 002** Dec 08, 2015**AB** PRINSTON INC **8MEQ** **A209026 001** Jun 11, 2019**AB** **10MEQ** **A209026 002** Jun 11, 2019**AB** TRIS PHARMA INC **8MEQ** **A201944 001** Mar 04, 2016**AB** **10MEQ** **A201944 002** Mar 04, 2016**AB** ZYDUS PHARMS **8MEQ** **A208445 001** Mar 11, 2019**AB** **10MEQ** **A208445 002** Mar 11, 2019

FOR SOLUTION; ORAL

KLOR-CON**AA** UPSHER SMITH LABS **20MEQ** **A209662 001** Oct 23, 2017POTASSIUM CHLORIDE**AA** AMNEAL PHARMS LLC **20MEQ** **A210902 001** May 23, 2019**AA** BELCHER **20MEQ** **A212183 001** May 06, 2019**AA** EPIC PHARMA LLC **20MEQ** **A210200 001** Nov 23, 2018**AA** NOVEL LABS INC **20MEQ** **A210241 001** Nov 21, 2018**AA** +! PHARMA RES SOFTWARE **20MEQ** **N208019 001** Aug 19, 2015

INJECTABLE; INJECTION

POTASSIUM CHLORIDE**AP** B BRAUN **2MEQ/ML** **A085870 001****AP** FRESENIUS KABI USA **2MEQ/ML** **A080225 001****AP** ! HOSPIRA **2MEQ/ML** **A080205 001**POTASSIUM CHLORIDE 10MEQ IN PLASTIC CONTAINER**AP** +! BAXTER HLTHCARE **14.9MG/ML** **N019904 001** Dec 26, 1989**AP** +! **746MG/100ML** **N019904 005** Dec 17, 1990**AP** + ICU MEDICAL INC **14.9MG/ML** **N020161 005** Nov 30, 1992**AP** + **745MG/100ML** **N020161 001** Nov 30, 1992

PRESCRIPTION DRUG PRODUCT LIST

POTASSIUM CHLORIDE

INJECTABLE; INJECTION

POTASSIUM CHLORIDE 20MEQ IN PLASTIC CONTAINER

<u>AP</u>	+	BAXTER HLTHCARE	<u>29.8MG/ML</u>	<u>N019904 002</u>	Dec 26, 1989
<u>AP</u>	+		<u>1.49GM/100ML</u>	<u>N019904 006</u>	Dec 17, 1990
<u>AP</u>	+	ICU MEDICAL INC	<u>29.8MG/ML</u>	<u>N020161 006</u>	Aug 11, 1998
<u>AP</u>	+		<u>1.49GM/100ML</u>	<u>N020161 002</u>	Nov 30, 1992

POTASSIUM CHLORIDE 40MEQ IN PLASTIC CONTAINER

<u>AP</u>	+	BAXTER HLTHCARE	<u>2.98GM/100ML</u>	<u>N019904 004</u>	Dec 26, 1989
<u>AP</u>	+	ICU MEDICAL INC	<u>2.98GM/100ML</u>	<u>N020161 004</u>	Aug 11, 1998

POTASSIUM CHLORIDE IN PLASTIC CONTAINER

<u>AP</u>		FRESENIUS KABI USA	<u>2MEQ/ML</u>	<u>A088901 001</u>	Jan 25, 1985
<u>AP</u>			<u>2MEQ/ML</u>	<u>A088908 001</u>	Jan 25, 1985

POTASSIUM CHLORIDE

! FRESENIUS KABI USA 3MEQ/ML A080225 003

POTASSIUM CHLORIDE 30MEQ IN PLASTIC CONTAINER

! BAXTER HLTHCARE 2.24GM/100ML N019904 003 Dec 26, 1989

SOLUTION; ORAL

POTASSIUM CHLORIDE

<u>AA</u>		AMNEAL PHARMS LLC	<u>20MEQ/15ML</u>	<u>A210041 001</u>	Jul 19, 2018
<u>AA</u>			<u>40MEQ/15ML</u>	<u>A210041 002</u>	Jul 19, 2018
<u>AA</u>		APOTEX	<u>20MEQ/15ML</u>	<u>A211067 001</u>	Aug 08, 2018
<u>AA</u>			<u>40MEQ/15ML</u>	<u>A211067 002</u>	Aug 08, 2018
<u>AA</u>	+	GENUS LIFESCIENCES	<u>20MEQ/15ML</u>	<u>N206814 001</u>	Dec 22, 2014
<u>AA</u>	+		<u>40MEQ/15ML</u>	<u>N206814 002</u>	Dec 22, 2014
<u>AA</u>		NOVEL LABS INC	<u>20MEQ/15ML</u>	<u>A209786 001</u>	Aug 29, 2018
<u>AA</u>			<u>40MEQ/15ML</u>	<u>A209786 002</u>	Aug 29, 2018
<u>AA</u>		PHARM ASSOC	<u>20MEQ/15ML</u>	<u>A210766 001</u>	Mar 29, 2019
<u>AA</u>			<u>40MEQ/15ML</u>	<u>A210766 002</u>	Mar 29, 2019

TABLET, EXTENDED RELEASE; ORAL

KLOR-CON M10

<u>AB1</u>		UPSHER SMITH LABS	<u>10MEQ</u>	<u>A074726 002</u>	Aug 09, 2000
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KLOR-CON M15

<u>AB1</u>		UPSHER SMITH LABS	<u>15MEQ</u>	<u>A074726 003</u>	Jun 06, 2003
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KLOR-CON M20

<u>AB1</u>	!	UPSHER SMITH LABS	<u>20MEQ</u>	<u>A074726 001</u>	Nov 20, 1998
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POTASSIUM CHLORIDE

<u>AB1</u>		ACTAVIS LABS FL INC	<u>10MEQ</u>	<u>A075604 001</u>	Apr 10, 2002
<u>AB1</u>			<u>20MEQ</u>	<u>A075604 002</u>	Apr 10, 2002
<u>AB1</u>		ADARE PHARMS INC	<u>10MEQ</u>	<u>A076368 002</u>	Jun 05, 2019
<u>AB1</u>			<u>20MEQ</u>	<u>A076368 001</u>	Aug 18, 2004
<u>AB1</u>		GLENMARK PHARMS LTD	<u>10MEQ</u>	<u>A203562 001</u>	Jul 26, 2016
<u>AB1</u>			<u>20MEQ</u>	<u>A203562 002</u>	Jul 26, 2016
<u>AB1</u>		NOVEL LABS INC	<u>10MEQ</u>	<u>A206347 001</u>	Jan 21, 2016
<u>AB1</u>			<u>20MEQ</u>	<u>A206347 002</u>	Jan 21, 2016
<u>AB1</u>		PRINSTON INC	<u>10MEQ</u>	<u>A209922 001</u>	Apr 30, 2019
<u>AB1</u>			<u>15MEQ</u>	<u>A209922 002</u>	Apr 30, 2019
<u>AB1</u>			<u>20MEQ</u>	<u>A209922 003</u>	Apr 30, 2019

KLOR-CON

<u>AB2</u>	+	UPSHER SMITH LABS	<u>8MEQ</u>	<u>N019123 001</u>	Apr 17, 1986
<u>AB2</u>	+		<u>10MEQ</u>	<u>N019123 002</u>	Apr 17, 1986

POTASSIUM CHLORIDE

<u>AB2</u>		AUROBINDO PHARMA LTD	<u>8MEQ</u>	<u>A210921 001</u>	Dec 19, 2018
<u>AB2</u>			<u>10MEQ</u>	<u>A210921 002</u>	Dec 19, 2018
<u>AB2</u>		GRANULES PHARMS	<u>8MEQ</u>	<u>A211797 001</u>	Mar 04, 2020
<u>AB2</u>			<u>10MEQ</u>	<u>A211797 002</u>	Mar 04, 2020
<u>AB2</u>		MYLAN	<u>8MEQ</u>	<u>A204662 001</u>	Aug 21, 2014
<u>AB2</u>			<u>10MEQ</u>	<u>A204662 002</u>	Aug 21, 2014
<u>AB2</u>		NOVEL LABS INC	<u>8MEQ</u>	<u>A206759 001</u>	Aug 09, 2016
<u>AB2</u>			<u>10MEQ</u>	<u>A206759 002</u>	Aug 09, 2016
<u>AB2</u>		PADDOCK LLC	<u>8MEQ</u>	<u>A205993 001</u>	Nov 05, 2015
<u>AB2</u>			<u>10MEQ</u>	<u>A205993 002</u>	Nov 05, 2015
<u>AB2</u>		SIGMAPHARM LABS LLC	<u>8MEQ</u>	<u>A207528 001</u>	Aug 19, 2016
<u>AB2</u>			<u>10MEQ</u>	<u>A207528 002</u>	Aug 19, 2016
<u>AB2</u>		STRIDES PHARMA	<u>8MEQ</u>	<u>A210733 001</u>	Aug 31, 2018
<u>AB2</u>			<u>10MEQ</u>	<u>A210733 002</u>	Aug 31, 2018
<u>AB2</u>		YICHANG HUMANWELL	<u>8MEQ</u>	<u>A209314 001</u>	Jun 22, 2018
<u>AB2</u>			<u>10MEQ</u>	<u>A209314 002</u>	Jun 22, 2018

K-TAB

<u>AB3</u>	+	ABBVIE	<u>10MEQ</u>	<u>N018279 001</u>	
<u>AB3</u>	+		<u>20MEQ</u>	<u>N018279 003</u>	Nov 25, 2013

PRESCRIPTION DRUG PRODUCT LISTPOTASSIUM CHLORIDE

TABLET, EXTENDED RELEASE;ORAL

POTASSIUM CHLORIDE

AB3	VITRUVIAS THERAP	10MEQ	A209688 001	Jan 12, 2018
AB3		20MEQ	A209688 002	Jan 12, 2018
AB3	YICHANG HUMANWELL	10MEQ	A212561 001	Sep 30, 2019
AB3		20MEQ	A212561 002	Sep 30, 2019
K-TAB				
BC	+ ABBVIE	8MEQ	N018279 002	Aug 01, 1988

POTASSIUM CHLORIDE; SODIUM CHLORIDE

INJECTABLE; INJECTION

POTASSIUM CHLORIDE 0.149% IN SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER

AP	ICU MEDICAL INC	149MG/100ML;450MG/100ML	A078446 001	Sep 10, 2008
<u>POTASSIUM CHLORIDE 0.15% IN SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER</u>				
AP	+! BAXTER HLTHCARE	150MG/100ML;450MG/100ML	N017648 005	Nov 26, 2002
<u>POTASSIUM CHLORIDE 0.15% IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER</u>				
AP	B BRAUN	150MG/100ML;900MG/100ML	N019708 004	Sep 29, 1989
AP	+ BAXTER HLTHCARE	150MG/100ML;900MG/100ML	N017648 001	
<u>POTASSIUM CHLORIDE 0.3% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER</u>				
AP	+ BAXTER HLTHCARE	300MG/100ML;900MG/100ML	N017648 002	
<u>POTASSIUM CHLORIDE 20MEQ IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER</u>				
AP	ICU MEDICAL INC	149MG/100ML;900MG/100ML	N019686 001	Oct 17, 1988
<u>POTASSIUM CHLORIDE 40MEQ IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER</u>				
AP	ICU MEDICAL INC	298MG/100ML;900MG/100ML	N019686 002	Oct 17, 1988

POTASSIUM CITRATE

TABLET, EXTENDED RELEASE;ORAL

POTASSIUM CITRATE

AB	ANI PHARMS INC	10MEQ	A212779 001	Jan 14, 2020
AB		15MEQ	A212779 002	Jan 14, 2020
AB	MOUNTAIN	5MEQ	A077440 001	Jun 09, 2006
AB		10MEQ	A077440 002	Jun 09, 2006
AB	STRIDES PHARMA	5MEQ	A206813 001	Sep 11, 2017
AB		10MEQ	A206813 002	Sep 11, 2017
AB		15MEQ	A206813 003	Sep 11, 2017
AB	TEVA PHARMS USA INC	5MEQ	A209758 001	Mar 05, 2018
AB		10MEQ	A209758 002	Mar 05, 2018
AB		15MEQ	A209758 003	Mar 05, 2018
AB	ZYDUS PHARMS	5MEQ	A203546 001	Aug 06, 2014
AB		10MEQ	A203546 002	Aug 06, 2014
AB		15MEQ	A203546 003	Aug 06, 2014

UROCI-T-K

AB	+ MISSION PHARMA	5MEQ	N019071 001	Aug 30, 1985
AB		10MEQ	N019071 002	Aug 31, 1992
AB	+!	15MEQ	N019071 003	Dec 30, 2009

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
+!	FRESENIUS KABI USA	1.18GM/5ML (236MG/ML);1.12GM/5ML (224MG/ML)	N212832 001	Nov 26, 2019
+!		3.54GM/15ML (236MG/ML);3.36GM/15ML (224MG/ML)	N212832 002	Nov 26, 2019
+!		11.8GM/50ML (236MG/ML);11.2GM/50ML (224MG/ML)	N212832 003	Nov 26, 2019

POVIDONE-IODINE

SOLUTION/DROPS;OPHTHALMIC

BETADINE

+!	ALCON PHARMS LTD	5%	N018634 001	Dec 17, 1986
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PRALATREXATE

SOLUTION; INTRAVENOUS

FOLOTYN

+	ACROTECH	20MG/ML (20MG/ML)	N022468 001	Sep 24, 2009
+!		40MG/2ML (20MG/ML)	N022468 002	Sep 24, 2009

PRESCRIPTION DRUG PRODUCT LIST

PRALIDOXIME CHLORIDE

INJECTABLE; INJECTION

PRALIDOXIME CHLORIDE

+! MERIDIAN MEDCL 300MG/ML

N018986 001 Apr 26, 1983

TECHN

PROTOPAM CHLORIDE

+! BAXTER HLTHCARE 1GM/VIAL

N014134 001

CORP

PRAMIPEXOLE DIHYDROCHLORIDE

TABLET; ORAL

MIRAPEX

AB + BOEHRINGER 0.125MG **N020667 001** Jul 01, 1997
INGELHEIM

AB +! 0.25MG **N020667 002** Jul 01, 1997

AB + 0.5MG **N020667 006** Feb 12, 1998

AB + 0.75MG **N020667 007** Jul 30, 2007

AB + 1MG **N020667 003** Jul 01, 1997

AB + 1.5MG **N020667 005** Jul 01, 1997

PRAMIPEXOLE DIHYDROCHLORIDE

AB ALEMBIC PHARMS LTD 0.125MG **A078894 001** Oct 08, 2010

AB 0.25MG **A078894 002** Oct 08, 2010

AB 0.5MG **A078894 003** Oct 08, 2010

AB 1MG **A078894 004** Oct 08, 2010

AB 1.5MG **A078894 005** Oct 08, 2010

AB AUROBINDO PHARMA 0.125MG **A202633 001** Oct 26, 2012
LTD

AB 0.25MG **A202633 002** Oct 26, 2012

AB 0.5MG **A202633 003** Oct 26, 2012

AB 0.75MG **A202633 004** Oct 26, 2012

AB 1MG **A202633 005** Oct 26, 2012

AB 1.5MG **A202633 006** Oct 26, 2012

AB BRECKENRIDGE PHARM 0.125MG **A091450 001** Oct 08, 2010

AB 0.25MG **A091450 002** Oct 08, 2010

AB 0.5MG **A091450 003** Oct 08, 2010

AB 1MG **A091450 004** Oct 08, 2010

AB 1.5MG **A091450 005** Oct 08, 2010

AB CSPC OUYI 0.25MG **A211088 001** Oct 03, 2018

AB 0.5MG **A211088 002** Oct 03, 2018

AB 0.75MG **A211088 003** Oct 03, 2018

AB 1MG **A211088 004** Oct 03, 2018

AB 1.5MG **A211088 005** Oct 03, 2018

AB GLENMARK GENERICS 0.125MG **A090781 001** Oct 08, 2010

AB 0.25MG **A090781 002** Oct 08, 2010

AB 0.5MG **A090781 003** Oct 08, 2010

AB 0.75MG **A090781 006** Sep 11, 2015

AB 1MG **A090781 004** Oct 08, 2010

AB 1.5MG **A090781 005** Oct 08, 2010

AB HERITAGE PHARMA 0.125MG **A077724 001** Feb 19, 2008

AB 0.25MG **A077724 002** Feb 19, 2008

AB 0.5MG **A077724 003** Feb 19, 2008

AB 1MG **A077724 004** Feb 19, 2008

AB 1.5MG **A077724 005** Feb 19, 2008

AB MACLEODS PHARMS LTD 0.125MG **A202164 001** Sep 20, 2012

AB 0.25MG **A202164 002** Sep 20, 2012

AB 0.5MG **A202164 003** Sep 20, 2012

AB 1MG **A202164 004** Sep 20, 2012

AB 1.5MG **A202164 005** Sep 20, 2012

AB SCIEGEN PHARMS INC 0.125MG **A203855 001** Oct 28, 2014

AB 0.25MG **A203855 002** Oct 28, 2014

AB 0.5MG **A203855 003** Oct 28, 2014

AB 0.75MG **A203855 004** Oct 28, 2014

AB 1MG **A203855 005** Oct 28, 2014

AB 1.5MG **A203855 006** Oct 28, 2014

AB STRIDES PHARMA 0.125MG **A202702 001** Jun 03, 2014

AB 0.25MG **A202702 002** Jun 03, 2014

AB 0.5MG **A202702 003** Jun 03, 2014

AB 0.75MG **A202702 004** Jun 03, 2014

AB 1MG **A202702 005** Jun 03, 2014

AB 1.5MG **A202702 006** Jun 03, 2014

AB TORRENT PHARMS 0.125MG **A090865 001** Oct 08, 2010

AB 0.25MG **A090865 002** Oct 08, 2010

AB 0.5MG **A090865 003** Oct 08, 2010

AB 0.75MG **A090865 004** Oct 08, 2010

PRESCRIPTION DRUG PRODUCT LIST

PRAMIPEXOLE DIHYDROCHLORIDE

TABLET;ORAL

PRAMIPEXOLE DIHYDROCHLORIDE

<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>	UNICHEM LABS LTD	<u>0.125MG</u>	<u>A207011 001</u>	Dec 19, 2018
<u>AB</u>		<u>0.25MG</u>	<u>A207011 002</u>	Dec 19, 2018
<u>AB</u>		<u>0.5MG</u>	<u>A207011 003</u>	Dec 19, 2018
<u>AB</u>		<u>0.75MG</u>	<u>A207011 004</u>	Dec 19, 2018
<u>AB</u>		<u>1MG</u>	<u>A207011 005</u>	Dec 19, 2018
<u>AB</u>		<u>1.5MG</u>	<u>A207011 006</u>	Dec 19, 2018
<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>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

MIRAPEX ER

<u>AB</u>	+!	BOEHRINGER INGELHEIM	<u>0.375MG</u>	<u>N022421 001</u>	Feb 19, 2010
<u>AB</u>	+		<u>0.75MG</u>	<u>N022421 002</u>	Feb 19, 2010
<u>AB</u>	+		<u>1.5MG</u>	<u>N022421 003</u>	Feb 19, 2010
<u>AB</u>	+		<u>2.25MG</u>	<u>N022421 006</u>	Jun 17, 2011
<u>AB</u>	+		<u>3MG</u>	<u>N022421 004</u>	Feb 19, 2010
<u>AB</u>	+		<u>3.75MG</u>	<u>N022421 007</u>	Jun 17, 2011
<u>AB</u>	+		<u>4.5MG</u>	<u>N022421 005</u>	Feb 19, 2010

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 PHARMS LTD	<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
<u>AB</u>		ANCHEN PHARMS	<u>0.375MG</u>	<u>A202206 001</u>	Feb 06, 2014
<u>AB</u>			<u>0.75MG</u>	<u>A202206 002</u>	Feb 06, 2014
<u>AB</u>			<u>1.5MG</u>	<u>A202206 003</u>	Feb 06, 2014
<u>AB</u>			<u>2.25MG</u>	<u>A202206 004</u>	Feb 06, 2014
<u>AB</u>			<u>3MG</u>	<u>A202206 005</u>	Feb 06, 2014
<u>AB</u>			<u>3.75MG</u>	<u>A202206 006</u>	Feb 06, 2014
<u>AB</u>			<u>4.5MG</u>	<u>A202206 007</u>	Feb 06, 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>		SANDOZ INC	<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>		ZYDUS PHARMS	<u>0.375MG</u>	<u>A202891 001</u>	Dec 12, 2017

PRESCRIPTION DRUG PRODUCT LIST

PRAMIPEXOLE DIHYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL

PRAMIPEXOLE DIHYDROCHLORIDE

<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

+	AMAG PHARMS INC	6.5MG	N208470 001	Nov 16, 2016
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PRASUGREL HYDROCHLORIDE

TABLET;ORAL

EFFIENT

<u>AB</u>	+	ELI LILLY AND CO	<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 LTD	<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 BIOTEC LTD	<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
<u>AB</u>		USPHARMA WINDLAS	<u>EQ 5MG BASE</u>	<u>A205790 001</u>	Oct 16, 2017
<u>AB</u>			<u>EQ 10MG BASE</u>	<u>A205790 002</u>	Oct 16, 2017

PRAVASTATIN SODIUM

TABLET;ORAL

PRAVACHOL

<u>AB</u>	+	BRISTOL MYERS SQUIBB	<u>20MG</u>	<u>N019898 003</u>	Oct 31, 1991
<u>AB</u>	+		<u>40MG</u>	<u>N019898 004</u>	Mar 22, 1993
<u>AB</u>	+	!	<u>80MG</u>	<u>N019898 008</u>	Dec 18, 2001

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 LTD	<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>		CHARTWELL RX	<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>		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

PRESCRIPTION DRUG PRODUCT LIST

PRAVASTATIN SODIUM

TABLET;ORAL

PRAVASTATIN SODIUM

<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 GENERICS	<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>	HISUN PHARM HANGZHOU	<u>20MG</u>	<u>A206061 001</u>	Nov 23, 2018
<u>AB</u>		<u>40MG</u>	<u>A206061 002</u>	Nov 23, 2018
<u>AB</u>		<u>80MG</u>	<u>A206061 003</u>	Nov 23, 2018
<u>AB</u>	LUPIN PHARMS	<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>	MYLAN PHARMS INC	<u>10MG</u>	<u>A079187 001</u>	May 27, 2010
<u>AB</u>		<u>20MG</u>	<u>A079187 002</u>	May 27, 2010
<u>AB</u>		<u>40MG</u>	<u>A079187 003</u>	May 27, 2010
<u>AB</u>		<u>80MG</u>	<u>A079187 004</u>	May 27, 2010
<u>AB</u>	TEVA	<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>	TEVA PHARMS	<u>80MG</u>	<u>A077793 001</u>	Jan 15, 2008
<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>10MG</u>	<u>A077751 001</u>	Apr 30, 2008
<u>AB</u>		<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

PRAZICUANTEL

TABLET;ORAL

BILTRICIDE

<u>AB</u>	+! BAYER HLTHCARE	<u>600MG</u>	<u>N018714 001</u>	Dec 29, 1982
<u>AB</u>	<u>PRAZICUANTEL</u> PAR PHARM INC	<u>600MG</u>	<u>A208820 001</u>	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>	
	<u>PRAZOSIN HYDROCHLORIDE</u>			
<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

CREAM;TOPICAL

DERMATOP E EMOLLIENT

<u>AB</u>	+! VALEANT BERMUDA	<u>0.1%</u>	<u>N020279 001</u>	Oct 29, 1993
<u>AB</u>	<u>PREDNICARBATE</u> FOUGERA PHARMS	<u>0.1%</u>	<u>A077287 001</u>	Sep 19, 2006
	OINTMENT;TOPICAL			
	PREDNICARBATE			
	! FOUGERA PHARMS	<u>0.1%</u>	<u>A077236 001</u>	Mar 09, 2007

PRESCRIPTION DRUG PRODUCT LIST

PREDNISOLONE

SYRUP; ORAL

PREDNISOLONE

AA	!	HI TECH PHARMA CO	<u>15MG/5ML</u>	<u>A040401</u>	<u>001</u>	Feb 27, 2003
AA		LANNETT CO INC	<u>15MG/5ML</u>	<u>A040775</u>	<u>001</u>	Sep 21, 2007
AA		PHARM ASSOC	<u>15MG/5ML</u>	<u>A040399</u>	<u>001</u>	Mar 05, 2003
AA		VISTAPHARM	<u>15MG/5ML</u>	<u>A040323</u>	<u>001</u>	May 13, 1999
AA		WOCKHARDT BIO AG	<u>15MG/5ML</u>	<u>A040313</u>	<u>001</u>	Sep 10, 2003

PRELONE

AA		TEVA	<u>15MG/5ML</u>	<u>A089081</u>	<u>001</u>	Feb 04, 1986
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TABLET; ORAL

PREDNISOLONE

! WATSON LABS

5MG

A080354 001

PREDNISOLONE ACETATE

SUSPENSION/DROPS; OPHTHALMIC

OMNIPRED

AB	+	!	NOVARTIS	<u>1%</u>	<u>N017469</u>	<u>001</u>
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PRED FORTE

AB	+	!	ALLERGAN	<u>1%</u>	<u>N017011</u>	<u>001</u>
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PRED MILD

+! ALLERGAN

0.12%

N017100 001

PREDNISOLONE ACETATE; SULFACETAMIDE SODIUM

OINTMENT; OPHTHALMIC

BLEPHAMIDE S.O.P.

! ALLERGAN

0.2%;10%

A087748 001 Dec 03, 1986

SUSPENSION; OPHTHALMIC

BLEPHAMIDE

+! ALLERGAN

0.2%;10%

N012813 002

PREDNISOLONE SODIUM PHOSPHATE

SOLUTION; ORAL

PEDIAAPRED

AA	+	!	SETON PHARM	<u>EQ 5MG BASE/5ML</u>	<u>N019157</u>	<u>001</u>	May 28, 1986
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PREDNISOLONE SODIUM PHOSPHATE

AA			CHARTWELL RX	<u>EQ 5MG BASE/5ML</u>	<u>A075988</u>	<u>001</u>	May 25, 2004
AA			EDENBRIDGE PHARMS	<u>EQ 10MG BASE/5ML</u>	<u>A203559</u>	<u>001</u>	Dec 20, 2016
AA				<u>EQ 20MG BASE/5ML</u>	<u>A203559</u>	<u>002</u>	Dec 20, 2016
AA			HI TECH PHARMA	<u>EQ 5MG BASE/5ML</u>	<u>A075183</u>	<u>001</u>	Mar 26, 2003
AA	!		PHARM ASSOC	<u>EQ 10MG BASE/5ML</u>	<u>A078465</u>	<u>001</u>	Mar 07, 2008
AA				<u>EQ 15MG BASE/5ML</u>	<u>A076913</u>	<u>001</u>	Apr 25, 2005
AA	!			<u>EQ 20MG BASE/5ML</u>	<u>A078988</u>	<u>001</u>	Jun 09, 2008
AA			VINTAGE	<u>EQ 15MG BASE/5ML</u>	<u>A079010</u>	<u>001</u>	May 26, 2009
AA			WOCKHARDT BIO AG	<u>EQ 5MG BASE/5ML</u>	<u>A075099</u>	<u>001</u>	Jun 28, 2002
AA	!			<u>EQ 15MG BASE/5ML</u>	<u>A076895</u>	<u>001</u>	Oct 04, 2004
	!		MISSION PHARMA	EQ 25MG BASE/5ML	A091396	001	Sep 13, 2010
			PHARM ASSOC	EQ 30MG BASE/5ML	A204962	001	Mar 11, 2020

SOLUTION/DROPS; OPHTHALMIC

PREDNISOLONE SODIUM PHOSPHATE

! BAUSCH AND LOMB

EQ 0.9% PHOSPHATE

A040070 001 Jul 29, 1994

TABLET, ORALLY DISINTEGRATING; ORAL

ORAPRED ODT

AB	+		CONCORDIA PHARMS	<u>EQ 10MG BASE</u>	<u>N021959</u>	<u>001</u>	Jun 01, 2006
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INC

AB	+			<u>EQ 15MG BASE</u>	<u>N021959</u>	<u>002</u>	Jun 01, 2006
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AB	+	!		<u>EQ 30MG BASE</u>	<u>N021959</u>	<u>003</u>	Jun 01, 2006
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PREDNISOLONE SODIUM PHOSPHATE

AB			MYLAN PHARMS INC	<u>EQ 10MG BASE</u>	<u>A202179</u>	<u>001</u>	Apr 10, 2013
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AB				<u>EQ 15MG BASE</u>	<u>A202179</u>	<u>002</u>	Apr 10, 2013
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AB				<u>EQ 30MG BASE</u>	<u>A202179</u>	<u>003</u>	Apr 10, 2013
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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

PREDNISONE

SOLUTION; ORAL

PREDNISONE

! HIKMA

5MG/5ML

A088703 001 Nov 08, 1984

PREDNISONE INTENSOL

! HIKMA

5MG/ML

A088810 001 Feb 20, 1985

PRESCRIPTION DRUG PRODUCT LIST

PREDNISON

TABLET; ORAL

PREDNISON

<u>AB</u>	GENEYORK PHARMS	<u>1MG</u>	<u>A211496 001</u>	Dec 28, 2018
<u>AB</u>		<u>2.5MG</u>	<u>A211495 001</u>	Dec 07, 2018
<u>AB</u>		<u>5MG</u>	<u>A211495 002</u>	Dec 07, 2018
<u>AB</u>		<u>10MG</u>	<u>A210525 001</u>	Dec 04, 2018
<u>AB</u>		<u>20MG</u>	<u>A210525 002</u>	Dec 04, 2018
<u>AB</u>		<u>50MG</u>	<u>A210525 003</u>	Dec 04, 2018
<u>AB</u>	! HIKMA	<u>1MG</u>	<u>A087800 001</u>	Apr 22, 1982
<u>AB</u>	!	<u>2.5MG</u>	<u>A087801 001</u>	Apr 22, 1982
<u>AB</u>	!	<u>5MG</u>	<u>A080352 001</u>	
<u>AB</u>	!	<u>10MG</u>	<u>A084122 001</u>	
<u>AB</u>	!	<u>20MG</u>	<u>A087342 001</u>	
<u>AB</u>	!	<u>50MG</u>	<u>A084283 001</u>	
<u>AB</u>	HIKMA PHARMS	<u>50MG</u>	<u>A088465 001</u>	Jun 01, 1984
<u>AB</u>	JUBILANT CADISTA	<u>1MG</u>	<u>A040611 001</u>	Jun 06, 2005
<u>AB</u>		<u>5MG</u>	<u>A040362 002</u>	Aug 29, 2001
<u>AB</u>		<u>10MG</u>	<u>A040362 001</u>	Aug 29, 2001
<u>AB</u>		<u>20MG</u>	<u>A040362 003</u>	Jun 29, 2005
<u>AB</u>	MYLAN	<u>5MG</u>	<u>A080292 001</u>	
<u>AB</u>		<u>10MG</u>	<u>A088832 001</u>	Dec 04, 1985
<u>AB</u>		<u>20MG</u>	<u>A083677 001</u>	
<u>AB</u>	NOVITIUM PHARMA	<u>2.5MG</u>	<u>A211575 001</u>	Nov 15, 2019
<u>AB</u>		<u>5MG</u>	<u>A211575 002</u>	Nov 15, 2019
<u>AB</u>		<u>10MG</u>	<u>A211575 003</u>	Nov 15, 2019
<u>AB</u>		<u>20MG</u>	<u>A211575 004</u>	Nov 15, 2019
<u>AB</u>		<u>50MG</u>	<u>A211575 005</u>	Nov 15, 2019
<u>AB</u>	SUN PHARM INDUSTRIES	<u>5MG</u>	<u>A089247 002</u>	Dec 04, 1985
<u>AB</u>		<u>10MG</u>	<u>A089247 003</u>	Dec 04, 1985
<u>AB</u>		<u>20MG</u>	<u>A089247 001</u>	Dec 04, 1985
<u>AB</u>	VINTAGE PHARMS	<u>1MG</u>	<u>A040584 001</u>	Dec 21, 2004
<u>AB</u>		<u>2.5MG</u>	<u>A040581 001</u>	Dec 21, 2004
<u>AB</u>		<u>5MG</u>	<u>A040256 001</u>	Jul 12, 2002
<u>AB</u>		<u>10MG</u>	<u>A040256 002</u>	Jul 12, 2002
<u>AB</u>		<u>20MG</u>	<u>A040392 001</u>	Feb 12, 2003
<u>AB</u>	WATSON LABS	<u>5MG</u>	<u>A080356 001</u>	
<u>AB</u>		<u>10MG</u>	<u>A085162 001</u>	
<u>AB</u>		<u>20MG</u>	<u>A085161 001</u>	

TABLET, DELAYED RELEASE; ORAL

PREDNISON

<u>AB</u>	ACTAVIS LABS FL INC	<u>1MG</u>	<u>A204867 001</u>	Apr 25, 2017
<u>AB</u>		<u>2MG</u>	<u>A204867 002</u>	Apr 25, 2017
<u>AB</u>		<u>5MG</u>	<u>A204867 003</u>	Apr 25, 2017
<u>RAYOS</u>				
<u>AB</u>	+ HORIZON PHARMA USA	<u>1MG</u>	<u>N202020 001</u>	Jul 26, 2012
<u>AB</u>	+	<u>2MG</u>	<u>N202020 002</u>	Jul 26, 2012
<u>AB</u>	+!	<u>5MG</u>	<u>N202020 003</u>	Jul 26, 2012

PREGABALIN

CAPSULE; ORAL

LYRICA

<u>AB</u>	+ PF PRISM CV	<u>25MG</u>	<u>N021446 001</u>	Dec 30, 2004
<u>AB</u>	+	<u>50MG</u>	<u>N021446 002</u>	Dec 30, 2004
<u>AB</u>	+	<u>75MG</u>	<u>N021446 003</u>	Dec 30, 2004
<u>AB</u>	+	<u>100MG</u>	<u>N021446 004</u>	Dec 30, 2004
<u>AB</u>	+	<u>150MG</u>	<u>N021446 005</u>	Dec 30, 2004
<u>AB</u>	+	<u>200MG</u>	<u>N021446 006</u>	Dec 30, 2004
<u>AB</u>	+	<u>225MG</u>	<u>N021446 007</u>	Dec 30, 2004
<u>AB</u>	+!	<u>300MG</u>	<u>N021446 008</u>	Dec 30, 2004

PREGABALIN

<u>AB</u>	ALEMBIC PHARMS LTD	<u>25MG</u>	<u>A203459 001</u>	Jul 19, 2019
<u>AB</u>		<u>50MG</u>	<u>A203459 002</u>	Jul 19, 2019
<u>AB</u>		<u>75MG</u>	<u>A203459 003</u>	Jul 19, 2019
<u>AB</u>		<u>100MG</u>	<u>A203459 004</u>	Jul 19, 2019
<u>AB</u>		<u>150MG</u>	<u>A203459 005</u>	Jul 19, 2019
<u>AB</u>		<u>200MG</u>	<u>A203459 006</u>	Jul 19, 2019
<u>AB</u>		<u>225MG</u>	<u>A203459 007</u>	Jul 19, 2019
<u>AB</u>		<u>300MG</u>	<u>A203459 008</u>	Jul 19, 2019
<u>AB</u>	ALKEM LABS LTD	<u>25MG</u>	<u>A211177 001</u>	Sep 30, 2019
<u>AB</u>		<u>50MG</u>	<u>A211177 002</u>	Sep 30, 2019
<u>AB</u>		<u>75MG</u>	<u>A207799 001</u>	Jul 19, 2019

PRESCRIPTION DRUG PRODUCT LIST

PREGABALIN

CAPSULE; ORAL

PREGABALIN

<u>AB</u>		<u>100MG</u>	<u>A207799 002</u>	Jul 19, 2019
<u>AB</u>		<u>150MG</u>	<u>A207799 003</u>	Jul 19, 2019
<u>AB</u>		<u>200MG</u>	<u>A207799 004</u>	Jul 19, 2019
<u>AB</u>		<u>225MG</u>	<u>A207799 005</u>	Jul 19, 2019
<u>AB</u>		<u>300MG</u>	<u>A207799 006</u>	Jul 19, 2019
<u>AB</u>	AMNEAL PHARMS CO	<u>25MG</u>	<u>A209743 001</u>	Jul 19, 2019
<u>AB</u>		<u>50MG</u>	<u>A209743 002</u>	Jul 19, 2019
<u>AB</u>		<u>75MG</u>	<u>A209743 003</u>	Jul 19, 2019
<u>AB</u>		<u>100MG</u>	<u>A209743 004</u>	Jul 19, 2019
<u>AB</u>		<u>150MG</u>	<u>A209743 005</u>	Jul 19, 2019
<u>AB</u>		<u>200MG</u>	<u>A209743 006</u>	Jul 19, 2019
<u>AB</u>		<u>225MG</u>	<u>A209743 007</u>	Jul 19, 2019
<u>AB</u>		<u>300MG</u>	<u>A209743 008</u>	Jul 19, 2019
<u>AB</u>	CIPLA	<u>25MG</u>	<u>A212280 001</u>	Jan 10, 2020
<u>AB</u>		<u>50MG</u>	<u>A212280 002</u>	Jan 10, 2020
<u>AB</u>		<u>75MG</u>	<u>A212280 003</u>	Jan 10, 2020
<u>AB</u>		<u>100MG</u>	<u>A212280 004</u>	Jan 10, 2020
<u>AB</u>		<u>150MG</u>	<u>A212280 005</u>	Jan 10, 2020
<u>AB</u>		<u>200MG</u>	<u>A212280 006</u>	Jan 10, 2020
<u>AB</u>		<u>225MG</u>	<u>A212280 007</u>	Jan 10, 2020
<u>AB</u>		<u>300MG</u>	<u>A212280 008</u>	Jan 10, 2020
<u>AB</u>	CSPC OUYI	<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>	DR REDDYS	<u>25MG</u>	<u>A209664 001</u>	Jul 19, 2019
<u>AB</u>		<u>50MG</u>	<u>A209664 002</u>	Jul 19, 2019
<u>AB</u>		<u>75MG</u>	<u>A209664 003</u>	Jul 19, 2019
<u>AB</u>		<u>100MG</u>	<u>A209664 004</u>	Jul 19, 2019
<u>AB</u>		<u>150MG</u>	<u>A209664 005</u>	Jul 19, 2019
<u>AB</u>		<u>200MG</u>	<u>A209664 006</u>	Jul 19, 2019
<u>AB</u>		<u>225MG</u>	<u>A209664 007</u>	Jul 19, 2019
<u>AB</u>		<u>300MG</u>	<u>A209664 008</u>	Jul 19, 2019
<u>AB</u>	HETERO LABS LTD III	<u>25MG</u>	<u>A206912 001</u>	Oct 08, 2019
<u>AB</u>		<u>50MG</u>	<u>A206912 002</u>	Oct 08, 2019
<u>AB</u>		<u>75MG</u>	<u>A206912 003</u>	Oct 08, 2019
<u>AB</u>		<u>100MG</u>	<u>A206912 004</u>	Oct 08, 2019
<u>AB</u>		<u>150MG</u>	<u>A206912 005</u>	Oct 08, 2019
<u>AB</u>		<u>200MG</u>	<u>A206912 006</u>	Oct 08, 2019
<u>AB</u>		<u>225MG</u>	<u>A206912 007</u>	Oct 08, 2019
<u>AB</u>		<u>300MG</u>	<u>A206912 008</u>	Oct 08, 2019
<u>AB</u>	INVAGEN PHARMS	<u>25MG</u>	<u>A211384 001</u>	Jul 19, 2019
<u>AB</u>		<u>50MG</u>	<u>A211384 002</u>	Jul 19, 2019
<u>AB</u>		<u>75MG</u>	<u>A211384 003</u>	Jul 19, 2019
<u>AB</u>		<u>100MG</u>	<u>A211384 004</u>	Jul 19, 2019
<u>AB</u>		<u>150MG</u>	<u>A211384 005</u>	Jul 19, 2019
<u>AB</u>		<u>200MG</u>	<u>A211384 006</u>	Jul 19, 2019
<u>AB</u>		<u>225MG</u>	<u>A211384 007</u>	Jul 19, 2019
<u>AB</u>		<u>300MG</u>	<u>A211384 008</u>	Jul 19, 2019
<u>AB</u>	MSN	<u>25MG</u>	<u>A209357 001</u>	Jul 19, 2019
<u>AB</u>		<u>50MG</u>	<u>A209357 002</u>	Jul 19, 2019
<u>AB</u>		<u>75MG</u>	<u>A209357 003</u>	Jul 19, 2019
<u>AB</u>		<u>100MG</u>	<u>A209357 004</u>	Jul 19, 2019
<u>AB</u>		<u>150MG</u>	<u>A209357 005</u>	Jul 19, 2019
<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>	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

PRESCRIPTION DRUG PRODUCT LIST

PREGABALIN

CAPSULE; ORAL

PREGABALIN

<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>	SHUANGCHENG	<u>75MG</u>	<u>A210891 001</u>	Sep 30, 2019
<u>AB</u>		<u>300MG</u>	<u>A210891 002</u>	Sep 30, 2019
<u>AB</u>	SUN PHARM	<u>25MG</u>	<u>A091157 001</u>	Nov 29, 2019
<u>AB</u>		<u>50MG</u>	<u>A091157 002</u>	Nov 29, 2019
<u>AB</u>		<u>75MG</u>	<u>A091157 003</u>	Nov 29, 2019
<u>AB</u>		<u>100MG</u>	<u>A091157 004</u>	Nov 29, 2019
<u>AB</u>		<u>150MG</u>	<u>A091157 005</u>	Nov 29, 2019
<u>AB</u>		<u>200MG</u>	<u>A091157 006</u>	Nov 29, 2019
<u>AB</u>		<u>225MG</u>	<u>A091157 007</u>	Nov 29, 2019
<u>AB</u>		<u>300MG</u>	<u>A091157 008</u>	Nov 29, 2019
<u>AB</u>	TEVA PHARMS	<u>25MG</u>	<u>A091219 001</u>	Jul 19, 2019
<u>AB</u>		<u>50MG</u>	<u>A091219 002</u>	Jul 19, 2019
<u>AB</u>		<u>75MG</u>	<u>A091224 001</u>	Jul 19, 2019
<u>AB</u>		<u>100MG</u>	<u>A091224 002</u>	Jul 19, 2019
<u>AB</u>		<u>150MG</u>	<u>A091224 003</u>	Jul 19, 2019
<u>AB</u>		<u>200MG</u>	<u>A091224 004</u>	Jul 19, 2019
<u>AB</u>		<u>225MG</u>	<u>A091224 005</u>	Jul 19, 2019
<u>AB</u>		<u>300MG</u>	<u>A091224 006</u>	Jul 19, 2019

SOLUTION; ORAL

LYRICA

<u>AA</u>	<u>+</u> !	PF PRISM CV	<u>20MG/ML</u>	<u>N022488 001</u>	Jan 04, 2010
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PREGABALIN

<u>AA</u>		ALKEM LABS LTD	<u>20MG/ML</u>	<u>A207623 001</u>	Jul 19, 2019
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TABLET, EXTENDED RELEASE; ORAL

LYRICA CR

	<u>+</u>	PF PRISM CV	82.5MG	N209501 001	Oct 11, 2017
	<u>+</u>		165MG	N209501 002	Oct 11, 2017
	<u>+</u> !		330MG	N209501 003	Oct 11, 2017

PRETOMANID

TABLET; ORAL

PRETOMANID

	<u>+</u> !	MYLAN IRELAND LTD	200MG	N212862 001	Aug 14, 2019
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PRILOCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

PRILOCAINE HYDROCHLORIDE

	<u>!</u>	SEPTODONT INC	4%	A079235 001	Sep 29, 2010
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PRIMAQUINE PHOSPHATE

TABLET; ORAL

PRIMAQUINE

<u>AB</u>	<u>+</u> !	SANOFI AVENTIS US	<u>EQ 15MG BASE</u>	<u>N008316 001</u>	
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PRIMAQUINE PHOSPHATE

<u>AB</u>		ALVOGEN INC	<u>EQ 15MG BASE</u>	<u>A203924 001</u>	Feb 03, 2014
<u>AB</u>		BAYSHORE PHARMS LLC	<u>EQ 15MG BASE</u>	<u>A204476 001</u>	Feb 25, 2014
<u>AB</u>		NOVAST LABS	<u>EQ 15MG BASE</u>	<u>A206043 001</u>	Jun 23, 2016

PRIMIDONE

TABLET; ORAL

MYSOLINE

<u>AB</u>	<u>+</u> !	VALEANT	<u>50MG</u>	<u>N009170 003</u>	
<u>AB</u>	<u>+</u>		<u>250MG</u>	<u>N009170 002</u>	

PRIMIDONE

<u>AB</u>		AMNEAL PHARM	<u>50MG</u>	<u>A040866 001</u>	Apr 23, 2008
<u>AB</u>			<u>250MG</u>	<u>A040866 002</u>	Apr 23, 2008
<u>AB</u>		ANDA REPOSITORY	<u>50MG</u>	<u>A040626 001</u>	Sep 29, 2005
<u>AB</u>			<u>250MG</u>	<u>A040626 002</u>	Sep 29, 2005
<u>AB</u>		HIKMA INTL PHARMS	<u>250MG</u>	<u>A040667 002</u>	Jul 27, 2006
<u>AB</u>		LANNETT	<u>50MG</u>	<u>A084903 002</u>	May 24, 2001
<u>AB</u>			<u>250MG</u>	<u>A084903 001</u>	
<u>AB</u>		OXFORD PHARMS	<u>50MG</u>	<u>A040586 001</u>	Feb 24, 2005
<u>AB</u>			<u>250MG</u>	<u>A040586 002</u>	Feb 24, 2005
<u>AB</u>		WATSON LABS	<u>250MG</u>	<u>A083551 001</u>	

PRESCRIPTION DRUG PRODUCT LIST

PROBENECID

TABLET; ORAL

PROBALAN

AB	LANNETT	500MG	A080966	001	
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PROBENECID

AB	MYLAN	500MG	A084211	002	
	! WATSON LABS TEVA	500MG	A084442	004	Mar 29, 1983

PROCAINAMIDE HYDROCHLORIDE

INJECTABLE; INJECTION

PROCAINAMIDE HYDROCHLORIDE

AP	! HOSPIRA	100MG/ML	A089069	001	Feb 12, 1986
AP	INTL MEDICATION	100MG/ML	A088636	001	Jul 31, 1984
AP	NEXUS PHARMS	100MG/ML	A206332	001	Oct 13, 2017
AP	! HOSPIRA	500MG/ML	A206332	002	Oct 13, 2017
		500MG/ML	A089070	001	Feb 12, 1986

PROCARBAZINE HYDROCHLORIDE

CAPSULE; ORAL

MATULANE

	+! LEADIANT BIOSCI INC	EQ 50MG BASE	N016785	001	
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PROCHLORPERAZINE

SUPPOSITORY; RECTAL

COMPRO

AB	PADDOCK LLC	25MG	A040246	001	Jun 28, 2000
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PROCHLORPERAZINE

AB	! ACP NIMBLE	25MG	A040058	001	Nov 24, 1993
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PROCHLORPERAZINE EDISYLATE

INJECTABLE; INJECTION

PROCHLORPERAZINE EDISYLATE

AP	ATHENEX INC	EQ 5MG BASE/ML	A040540	001	May 28, 2004
AP	! EMCURE PHARMS LTD	EQ 5MG BASE/ML	A204147	001	Oct 15, 2013
AP	MYLAN LABS LTD	EQ 5MG BASE/ML	A210710	001	Oct 25, 2018
AP	NEXUS PHARMS	EQ 5MG BASE/ML	A204860	001	Feb 15, 2019
AP	WEST-WARD PHARMS	EQ 5MG BASE/ML	A089903	001	Aug 29, 1989
	INT				

PROCHLORPERAZINE MALEATE

TABLET; ORAL

PROCHLORPERAZINE MALEATE

AB	MYLAN	EQ 5MG BASE	A040185	002	Oct 28, 1996
AB		EQ 10MG BASE	A040185	001	Oct 28, 1996
AB	SANDOZ	EQ 5MG BASE	A040101	001	Jul 19, 1996
AB	!	EQ 10MG BASE	A040101	002	Jul 19, 1996

PROCOMP

AB	JUBILANT CADISTA	EQ 5MG BASE	A040268	001	Feb 27, 1998
AB		EQ 10MG BASE	A040268	002	Feb 27, 1998

PROGESTERONE

CAPSULE; ORAL

PROGESTERONE

AB	AMNEAL PHARMS NY	100MG	A207724	001	Sep 07, 2017
AB		200MG	A207724	002	Sep 07, 2017
AB	BIONPHARMA INC	100MG	A200900	001	Aug 16, 2013
AB		200MG	A200900	002	Aug 16, 2013
AB	DR REDDYS	100MG	A208801	001	Feb 28, 2017
AB		200MG	A208801	002	Feb 28, 2017
AB	EUGIA PHARMA	100MG	A211285	001	Oct 26, 2018
AB		200MG	A211285	002	Oct 26, 2018
AB	SOFGEN PHARMS	100MG	A200456	001	Sep 28, 2012
AB		200MG	A200456	002	Sep 28, 2012
AB	XIROMED	100MG	A205229	001	Oct 20, 2017
AB		200MG	A205229	002	Oct 20, 2017

PROMETRIUM

AB	+ VIRTUS PHARMS	100MG	N019781	001	May 14, 1998
AB	+!	200MG	N019781	002	Oct 15, 1999

GEL; VAGINAL

CRINONE

	+! ALLERGAN	4%	N020701	001	Jul 31, 1997
	+!	8%	N020701	002	Jul 31, 1997

INJECTABLE; INJECTION

PROGESTERONE

AO	+! ACTAVIS LABS UT INC	50MG/ML	N017362	002	
AO	EUGIA PHARMA	50MG/ML	A210965	001	Dec 06, 2018

PRESCRIPTION DRUG PRODUCT LIST

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PROGESTERONE

INJECTABLE; INJECTION

PROGESTERONE

<u>AO</u>	FRESENIUS KABI USA	<u>50MG/ML</u>	<u>A075906</u>	<u>001</u>	Apr 25, 2001
<u>AO</u>	HIKMA FARMACEUTICA	<u>50MG/ML</u>	<u>A091033</u>	<u>001</u>	Oct 28, 2010

INSERT; VAGINAL

ENDOMETRIN

+	!	FERRING	100MG	N022057	001	Jun 21, 2007
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PROMETHAZINE HYDROCHLORIDE

INJECTABLE; INJECTION

PROMETHAZINE HYDROCHLORIDE

<u>AP</u>	!	WEST-WARD PHARMS	<u>25MG/ML</u>	<u>A083312</u>	<u>001</u>	
		INT				
<u>AP</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>		ACP NIMBLE	<u>12.5MG</u>	<u>A040428</u>	<u>002</u>	Mar 31, 2003
<u>AB</u>	!		<u>25MG</u>	<u>A040428</u>	<u>001</u>	Feb 05, 2002
<u>AB</u>		PERRIGO 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
<u>AB</u>		WATSON LABS INC	<u>12.5MG</u>	<u>A040479</u>	<u>001</u>	Jun 24, 2003
<u>AB</u>			<u>25MG</u>	<u>A040479</u>	<u>002</u>	Jun 24, 2003
		PROMETHEGAN				
	!	ACP NIMBLE	50MG	A087165	001	Aug 14, 1987

SYRUP; ORAL

PROMETHAZINE HYDROCHLORIDE

<u>AA</u>		AMNEAL PHARMS	<u>6.25MG/5ML</u>	<u>A040882</u>	<u>001</u>	Dec 30, 2009
<u>AA</u>		HI TECH PHARMA	<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>		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
<u>AA</u>		VINTAGE	<u>6.25MG/5ML</u>	<u>A040643</u>	<u>001</u>	Apr 26, 2006

PROMETHAZINE PLAIN

<u>AA</u>	!	WOCKHARDT BIO AG	<u>6.25MG/5ML</u>	<u>A087953</u>	<u>001</u>	Nov 15, 1982
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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>		SANDOZ	<u>25MG</u>	<u>A084176</u>	<u>003</u>	
<u>AB</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>		SUN PHARM INDS INC	<u>12.5MG</u>	<u>A040863</u>	<u>001</u>	Dec 30, 2008
<u>AB</u>			<u>25MG</u>	<u>A040863</u>	<u>002</u>	Dec 30, 2008
<u>AB</u>			<u>50MG</u>	<u>A040863</u>	<u>003</u>	Dec 30, 2008
<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>		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>		MYLAN	<u>225MG</u>	<u>A203803</u>	<u>001</u>	Apr 29, 2016

PRESCRIPTION DRUG PRODUCT LIST

PROPAFENONE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE;ORAL

PROPAFENONE HYDROCHLORIDE

<u>AB</u>		<u>325MG</u>	<u>A203803 002</u>	Apr 29, 2016
<u>AB</u>		<u>425MG</u>	<u>A203803 003</u>	Apr 29, 2016
<u>AB</u>	PAR PHARM	<u>225MG</u>	<u>A078540 001</u>	Oct 18, 2010
<u>AB</u>		<u>325MG</u>	<u>A078540 002</u>	Oct 18, 2010
<u>AB</u>		<u>425MG</u>	<u>A078540 003</u>	Oct 18, 2010
<u>AB</u>	SINOTHERAPEUTICS INC	<u>225MG</u>	<u>A210339 001</u>	Jan 04, 2019
<u>AB</u>		<u>325MG</u>	<u>A210339 002</u>	Jan 04, 2019
<u>AB</u>		<u>425MG</u>	<u>A210339 003</u>	Jan 04, 2019
<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>	WILSHIRE PHARMS INC	<u>225MG</u>	<u>A205956 001</u>	Jul 02, 2018
<u>AB</u>		<u>325MG</u>	<u>A205956 002</u>	Jul 02, 2018
<u>AB</u>		<u>425MG</u>	<u>A205956 003</u>	Jul 02, 2018
<u>RYTHMOL SR</u>				
<u>AB</u>	+ GLAXOSMITHKLINE LLC	<u>225MG</u>	<u>N021416 001</u>	Sep 04, 2003
<u>AB</u>	+	<u>325MG</u>	<u>N021416 002</u>	Sep 04, 2003
<u>AB</u>	+	<u>425MG</u>	<u>N021416 003</u>	Sep 04, 2003

TABLET; ORAL

PROPAFENONE HYDROCHLORIDE

<u>AB</u>	ANI PHARMS INC	<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 LTD	<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>	SUN PHARM INDUSTRIES	<u>150MG</u>	<u>A075998 001</u>	Nov 29, 2001
<u>AB</u>		<u>225MG</u>	<u>A075998 002</u>	Nov 29, 2001
<u>AB</u>		<u>300MG</u>	<u>A075998 003</u>	Nov 29, 2001
<u>AB</u>	VINTAGE PHARMS	<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

PROPANTHELINE BROMIDE

TABLET; ORAL

PROPANTHELINE BROMIDE

!	HIKMA	15MG	A080927 002	
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PROPARACAINE HYDROCHLORIDE

SOLUTION/DROPS;OPHTHALMIC

ALCAINE

<u>AT</u>	ALCON LABS INC	<u>0.5%</u>	<u>A080027 001</u>	
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PROPARACAINE HYDROCHLORIDE

<u>AT</u>	!	AKORN INC	<u>0.5%</u>	<u>A040277 001</u>	Mar 16, 2000
<u>AT</u>		BAUSCH AND LOMB	<u>0.5%</u>	<u>A040074 001</u>	Sep 29, 1995

PROPOFOL

INJECTABLE; INJECTION

DIPRIVAN

<u>AB</u>	+	FRESENIUS KABI USA	<u>10MG/ML</u>	<u>N019627 002</u>	Jun 11, 1996
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PROPOFOL

<u>AB</u>		DR REDDYS	<u>10MG/ML</u>	<u>A205067 001</u>	Nov 15, 2018
<u>AB</u>		HOSPIRA	<u>10MG/ML</u>	<u>A077908 001</u>	Mar 17, 2006
<u>AB</u>		SAGENT PHARMS INC	<u>10MG/ML</u>	<u>A075102 001</u>	Jan 04, 1999
<u>AB</u>		WATSON LABS INC	<u>10MG/ML</u>	<u>A205307 001</u>	Dec 22, 2015

PROPRANOLOL HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE;ORAL

INDERAL LA

<u>AB</u>	+	ANI PHARMS INC	<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

PRESCRIPTION DRUG PRODUCT LIST

3-370 (of 453)

PROPRANOLOL HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

PROPRANOLOL HYDROCHLORIDE

<u>AB</u>		<u>160MG</u>	<u>A078494</u>	<u>004</u>	Aug 10, 2007
<u>AB</u>	ADARE PHARMS INC	<u>60MG</u>	<u>A078703</u>	<u>001</u>	Jul 15, 2011
<u>AB</u>		<u>80MG</u>	<u>A078703</u>	<u>002</u>	Jul 15, 2011
<u>AB</u>		<u>120MG</u>	<u>A078703</u>	<u>003</u>	Jul 15, 2011
<u>AB</u>		<u>160MG</u>	<u>A078703</u>	<u>004</u>	Jul 15, 2011
<u>AB</u>	AMTA	<u>60MG</u>	<u>A212026</u>	<u>001</u>	Jan 06, 2020
<u>AB</u>		<u>80MG</u>	<u>A212026</u>	<u>002</u>	Jan 06, 2020
<u>AB</u>		<u>120MG</u>	<u>A212026</u>	<u>003</u>	Jan 06, 2020
<u>AB</u>		<u>160MG</u>	<u>A212026</u>	<u>004</u>	Jan 06, 2020
<u>AB</u>	NORTEC DEV ASSOC	<u>60MG</u>	<u>A078065</u>	<u>001</u>	Jan 26, 2007
<u>AB</u>		<u>80MG</u>	<u>A078065</u>	<u>002</u>	Jan 26, 2007
<u>AB</u>		<u>120MG</u>	<u>A078065</u>	<u>003</u>	Jan 26, 2007
<u>AB</u>		<u>160MG</u>	<u>A078065</u>	<u>004</u>	Jan 26, 2007
<u>AB</u>	ZYDUS PHARMS USA INC	<u>60MG</u>	<u>A090321</u>	<u>001</u>	Mar 25, 2011
<u>AB</u>		<u>80MG</u>	<u>A090321</u>	<u>002</u>	Mar 25, 2011
<u>AB</u>		<u>120MG</u>	<u>A090321</u>	<u>003</u>	Mar 25, 2011
<u>AB</u>		<u>160MG</u>	<u>A090321</u>	<u>004</u>	Mar 25, 2011
	INNOPRAN XL				
BX	+ ANI PHARMS INC	80MG	N021438	001	Mar 12, 2003
BX	+!	120MG	N021438	002	Mar 12, 2003

INJECTABLE; INJECTION

PROPRANOLOL HYDROCHLORIDE

<u>AP</u>	ATHENEX INC	<u>1MG/ML</u>	<u>A075792</u>	<u>001</u>	Aug 29, 2000
<u>AP</u>	FRESENIUS KABI USA	<u>1MG/ML</u>	<u>A075826</u>	<u>001</u>	Aug 31, 2001
	! HIKMA FARMACEUTICA	1MG/ML	A077760	001	Jan 31, 2008
	SOLUTION; ORAL				
	HEMANGEOL				
	+! PIERRE FABRE DERMA	4.28MG/ML	N205410	001	Mar 14, 2014
	PROPRANOLOL HYDROCHLORIDE				
	! HIKMA	20MG/5ML	A070979	001	May 15, 1987
	!	40MG/5ML	A070690	001	May 15, 1987

TABLET; ORAL

PROPRANOLOL HYDROCHLORIDE

<u>AB</u>	IMPAX LABS INC	<u>10MG</u>	<u>A071972</u>	<u>001</u>	Apr 06, 1988
<u>AB</u>		<u>20MG</u>	<u>A071972</u>	<u>002</u>	Apr 06, 1988
<u>AB</u>		<u>40MG</u>	<u>A071972</u>	<u>003</u>	Apr 06, 1988
<u>AB</u>		<u>60MG</u>	<u>A071976</u>	<u>002</u>	May 13, 1986
<u>AB</u>	!	<u>80MG</u>	<u>A071976</u>	<u>001</u>	Apr 06, 1988
<u>AB</u>	INNOGENIX	<u>10MG</u>	<u>A070322</u>	<u>002</u>	Oct 22, 1985
<u>AB</u>		<u>20MG</u>	<u>A070322</u>	<u>003</u>	Oct 22, 1985
<u>AB</u>		<u>40MG</u>	<u>A070322</u>	<u>004</u>	Oct 22, 1985
<u>AB</u>		<u>60MG</u>	<u>A070322</u>	<u>005</u>	Sep 24, 1986
<u>AB</u>		<u>80MG</u>	<u>A070322</u>	<u>001</u>	Aug 04, 1986
<u>AB</u>	IPCA LABS LTD	<u>10MG</u>	<u>A078955</u>	<u>001</u>	Jun 02, 2008
<u>AB</u>		<u>20MG</u>	<u>A078955</u>	<u>002</u>	Jun 02, 2008
<u>AB</u>		<u>40MG</u>	<u>A078955</u>	<u>003</u>	Jun 02, 2008
<u>AB</u>		<u>60MG</u>	<u>A078955</u>	<u>004</u>	Jun 02, 2008
<u>AB</u>		<u>80MG</u>	<u>A078955</u>	<u>005</u>	Jun 02, 2008
<u>AB</u>	MYLAN	<u>10MG</u>	<u>A070213</u>	<u>002</u>	Nov 19, 1985
<u>AB</u>		<u>20MG</u>	<u>A070213</u>	<u>003</u>	Nov 19, 1985
<u>AB</u>		<u>40MG</u>	<u>A070213</u>	<u>001</u>	Nov 19, 1985
<u>AB</u>		<u>60MG</u>	<u>A070213</u>	<u>005</u>	Apr 08, 2011
<u>AB</u>		<u>80MG</u>	<u>A070213</u>	<u>004</u>	Nov 19, 1985
<u>AB</u>	NORTHSTAR HLTHCARE	<u>10MG</u>	<u>A078213</u>	<u>001</u>	Jan 10, 2008
<u>AB</u>		<u>20MG</u>	<u>A078213</u>	<u>002</u>	Jan 10, 2008
<u>AB</u>		<u>40MG</u>	<u>A078213</u>	<u>003</u>	Jan 10, 2008
<u>AB</u>		<u>60MG</u>	<u>A078213</u>	<u>004</u>	Jan 10, 2008
<u>AB</u>		<u>80MG</u>	<u>A078213</u>	<u>005</u>	Jan 10, 2008
<u>AB</u>	VINTAGE PHARMS	<u>10MG</u>	<u>A070221</u>	<u>002</u>	Aug 01, 1986
<u>AB</u>		<u>20MG</u>	<u>A070221</u>	<u>003</u>	Aug 01, 1986
<u>AB</u>		<u>40MG</u>	<u>A070219</u>	<u>001</u>	Aug 01, 1986
<u>AB</u>		<u>40MG</u>	<u>A070221</u>	<u>004</u>	Aug 01, 1986
<u>AB</u>		<u>60MG</u>	<u>A070221</u>	<u>005</u>	Sep 24, 1986
<u>AB</u>		<u>80MG</u>	<u>A070221</u>	<u>001</u>	Apr 14, 1986
<u>AB</u>	WATSON LABS	<u>10MG</u>	<u>A070175</u>	<u>001</u>	May 13, 1986
<u>AB</u>		<u>20MG</u>	<u>A070176</u>	<u>001</u>	May 13, 1986
<u>AB</u>		<u>40MG</u>	<u>A070177</u>	<u>001</u>	May 13, 1986
<u>AB</u>		<u>60MG</u>	<u>A070178</u>	<u>002</u>	Apr 23, 2018
<u>AB</u>		<u>80MG</u>	<u>A070178</u>	<u>001</u>	May 13, 1986

PRESCRIPTION DRUG PRODUCT LISTPROPYLTHIOURACIL

TABLET; ORAL

PROPYLTHIOURACIL

BD	ACTAVIS ELIZABETH	50MG	A080172	001	
BD	+! DAVA PHARMS INC	50MG	N006188	001	

PROTAMINE SULFATE

INJECTABLE; INJECTION

PROTAMINE SULFATE

!	FRESENIUS KABI USA	10MG/ML	A089454	001	Apr 07, 1987
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PROTRIPTYLINE HYDROCHLORIDE

TABLET; ORAL

PROTRIPTYLINE HYDROCHLORIDE

AB	EPIC PHARMA LLC	5MG	A202220	001	Nov 19, 2012
AB		10MG	A202220	002	Nov 19, 2012
AB	HIKMA	5MG	A078913	001	Sep 16, 2008
AB		10MG	A078913	002	Sep 16, 2008
AB	SIGMAPHARM LABS LLC	5MG	A090462	001	May 03, 2010
AB		10MG	A090462	002	May 03, 2010
<u>VIVACTIL</u>					
AB	HERITAGE PHARMA	5MG	A073644	001	Aug 24, 1995
AB	!	10MG	A073645	001	Aug 24, 1995

PRUCALOPRIDE SUCCINATE

TABLET; ORAL

MOTEGRITY

+	SHIRE DEV LLC	EQ 1MG BASE	N210166	001	Dec 14, 2018
+	!	EQ 2MG BASE	N210166	002	Dec 14, 2018

PYRAZINAMIDE

TABLET; ORAL

PYRAZINAMIDE

!	AKORN	500MG	A081319	001	Jun 30, 1992
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PYRIDOSTIGMINE BROMIDE

INJECTABLE; INJECTION

MESTINON

AP	+! BAUSCH	5MG/ML	N009830	001	
<u>REGONOL</u>					
AP	SANDOZ INC	5MG/ML	N017398	001	

SYRUP; ORAL

MESTINON

AA	+! BAUSCH	60MG/5ML	N015193	001	
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PYRIDOSTIGMINE BROMIDE

AA	AMNEAL PHARMS LLC	60MG/5ML	A212702	001	Jan 10, 2020
AA	INVATECH	60MG/5ML	A208797	001	Jan 09, 2020
AA	NOVITIUM PHARMA	60MG/5ML	A211694	001	Mar 08, 2019

TABLET; ORAL

MESTINON

AB	+! BAUSCH	60MG	N009829	002	
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PYRIDOSTIGMINE BROMIDE

AB	ELYSIUM	60MG	A211181	001	Jul 20, 2018
AB	IMPAX LABS	60MG	A040502	001	Apr 24, 2003
AB	ZYDUS PHARMS	60MG	A205650	001	Jun 22, 2015
	ELYSIUM	30MG	A211181	002	May 14, 2019

TABLET, EXTENDED RELEASE; ORAL

MESTINON

AB	+! BAUSCH	180MG	N011665	001	
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PYRIDOSTIGMINE BROMIDE

AB	ALVOGEN	180MG	A204737	001	Jun 26, 2015
AB	IMPAX LABS INC	180MG	A203184	001	Sep 15, 2015
AB	RISING	180MG	A205464	001	Aug 15, 2017

PYRIDOXINE HYDROCHLORIDE

INJECTABLE; INJECTION

PYRIDOXINE HYDROCHLORIDE

!	FRESENIUS KABI USA	100MG/ML	A080618	001	
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PYRIMETHAMINE

TABLET; ORAL

DARAPRIM

AB	+! VYERA PHARMS LLC	25MG	N008578	001	
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PYRIMETHAMINE

AB	CEROVENE INC	25MG	A207127	001	Feb 28, 2020
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PRESCRIPTION DRUG PRODUCT LIST

QUAZEPAM

TABLET;ORAL

DORAL

+! GALT PHARMS

15MG

N018708 001 Dec 27, 1985

QUETIAPINE FUMARATE

TABLET;ORAL

QUETIAPINE FUMARATE

AB	ACCORD HLTHCARE	<u>EQ 25MG BASE</u>	<u>A202152 001</u>	Mar 27, 2012
AB		<u>EQ 50MG BASE</u>	<u>A202152 002</u>	Mar 27, 2012
AB		<u>EQ 100MG BASE</u>	<u>A202152 003</u>	Mar 27, 2012
AB		<u>EQ 200MG BASE</u>	<u>A202152 004</u>	Mar 27, 2012
AB		<u>EQ 300MG BASE</u>	<u>A202152 005</u>	Mar 27, 2012
AB		<u>EQ 400MG BASE</u>	<u>A202152 006</u>	Mar 27, 2012
AB	ALEMBIC PHARMS LTD	<u>EQ 25MG BASE</u>	<u>A203390 001</u>	Oct 28, 2014
AB		<u>EQ 50MG BASE</u>	<u>A203390 002</u>	Oct 28, 2014
AB		<u>EQ 100MG BASE</u>	<u>A203390 003</u>	Oct 28, 2014
AB		<u>EQ 200MG BASE</u>	<u>A203390 004</u>	Oct 28, 2014
AB		<u>EQ 300MG BASE</u>	<u>A203390 005</u>	Oct 28, 2014
AB		<u>EQ 400MG BASE</u>	<u>A203390 006</u>	Oct 28, 2014
AB	ALKEM LABS LTD	<u>EQ 25MG BASE</u>	<u>A201504 001</u>	Feb 12, 2013
AB		<u>EQ 50MG BASE</u>	<u>A201504 002</u>	Feb 12, 2013
AB		<u>EQ 100MG BASE</u>	<u>A201504 003</u>	Feb 12, 2013
AB		<u>EQ 150MG BASE</u>	<u>A201504 004</u>	Feb 12, 2013
AB		<u>EQ 200MG BASE</u>	<u>A201504 005</u>	Feb 12, 2013
AB		<u>EQ 300MG BASE</u>	<u>A201504 006</u>	Feb 12, 2013
AB		<u>EQ 400MG BASE</u>	<u>A201504 007</u>	Feb 12, 2013
AB	AUROBINDO PHARMA LTD	<u>EQ 25MG BASE</u>	<u>A091388 001</u>	Mar 27, 2012
AB		<u>EQ 50MG BASE</u>	<u>A091388 002</u>	Mar 27, 2012
AB		<u>EQ 100MG BASE</u>	<u>A091388 003</u>	Mar 27, 2012
AB		<u>EQ 150MG BASE</u>	<u>A091388 004</u>	Mar 27, 2012
AB		<u>EQ 200MG BASE</u>	<u>A091388 005</u>	Mar 27, 2012
AB		<u>EQ 300MG BASE</u>	<u>A091388 006</u>	Mar 27, 2012
AB		<u>EQ 400MG BASE</u>	<u>A091388 007</u>	Mar 27, 2012
AB	DR REDDYS LABS LTD	<u>EQ 25MG BASE</u>	<u>A077380 001</u>	Mar 27, 2012
AB		<u>EQ 50MG BASE</u>	<u>A077380 002</u>	Mar 27, 2012
AB		<u>EQ 100MG BASE</u>	<u>A077380 003</u>	Mar 27, 2012
AB		<u>EQ 150MG BASE</u>	<u>A077380 004</u>	Mar 27, 2012
AB		<u>EQ 200MG BASE</u>	<u>A077380 005</u>	Mar 27, 2012
AB		<u>EQ 300MG BASE</u>	<u>A077380 006</u>	Mar 27, 2012
AB		<u>EQ 400MG BASE</u>	<u>A077380 007</u>	Mar 27, 2012
AB	HIKMA	<u>EQ 25MG BASE</u>	<u>A090120 001</u>	Mar 27, 2012
AB		<u>EQ 50MG BASE</u>	<u>A090749 001</u>	Mar 27, 2012
AB		<u>EQ 100MG BASE</u>	<u>A090749 002</u>	Mar 27, 2012
AB		<u>EQ 200MG BASE</u>	<u>A090749 003</u>	Mar 27, 2012
AB		<u>EQ 300MG BASE</u>	<u>A090749 004</u>	Mar 27, 2012
AB		<u>EQ 400MG BASE</u>	<u>A090749 005</u>	Mar 27, 2012
AB	JUBILANT GENERICS	<u>EQ 25MG BASE</u>	<u>A203150 001</u>	Nov 26, 2013
AB	LUPIN LTD	<u>EQ 25MG BASE</u>	<u>A201109 001</u>	Mar 27, 2012
AB		<u>EQ 50MG BASE</u>	<u>A201109 002</u>	Mar 27, 2012
AB		<u>EQ 100MG BASE</u>	<u>A201109 003</u>	Mar 27, 2012
AB		<u>EQ 200MG BASE</u>	<u>A201109 004</u>	Mar 27, 2012
AB		<u>EQ 300MG BASE</u>	<u>A201109 005</u>	Mar 27, 2012
AB		<u>EQ 400MG BASE</u>	<u>A201109 006</u>	Mar 27, 2012
AB	MACLEODS PHARMS LTD	<u>EQ 25MG BASE</u>	<u>A203359 001</u>	May 17, 2016
AB		<u>EQ 50MG BASE</u>	<u>A203359 002</u>	May 17, 2016
AB		<u>EQ 100MG BASE</u>	<u>A203359 003</u>	May 17, 2016
AB		<u>EQ 200MG BASE</u>	<u>A203359 004</u>	May 17, 2016
AB		<u>EQ 300MG BASE</u>	<u>A203359 005</u>	May 17, 2016
AB		<u>EQ 400MG BASE</u>	<u>A203359 006</u>	May 17, 2016
AB	SANDOZ	<u>EQ 25MG BASE</u>	<u>A078679 001</u>	Dec 14, 2012
AB		<u>EQ 50MG BASE</u>	<u>A078679 002</u>	Dec 14, 2012
AB		<u>EQ 100MG BASE</u>	<u>A078679 003</u>	Dec 14, 2012
AB		<u>EQ 150MG BASE</u>	<u>A078679 004</u>	Dec 14, 2012
AB		<u>EQ 200MG BASE</u>	<u>A078679 005</u>	Dec 14, 2012
AB		<u>EQ 300MG BASE</u>	<u>A078679 006</u>	Dec 14, 2012
AB		<u>EQ 400MG BASE</u>	<u>A078679 007</u>	Dec 14, 2012
AB	SUN PHARM	<u>EQ 25MG BASE</u>	<u>A201190 001</u>	Mar 27, 2012
AB		<u>EQ 50MG BASE</u>	<u>A201190 002</u>	Mar 27, 2012
AB		<u>EQ 100MG BASE</u>	<u>A201190 003</u>	Mar 27, 2012
AB		<u>EQ 200MG BASE</u>	<u>A201190 004</u>	Mar 27, 2012
AB		<u>EQ 300MG BASE</u>	<u>A201190 005</u>	Mar 27, 2012

PRESCRIPTION DRUG PRODUCT LIST

QUETIAPINE FUMARATE

TABLET;ORAL

QUETIAPINE FUMARATE

<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>	TORRENT PHARMS LTD	<u>EQ 25MG BASE</u>	<u>A200363 001</u>	Mar 27, 2012
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A200363 002</u>	Mar 27, 2012
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A200363 003</u>	Mar 27, 2012
<u>AB</u>		<u>EQ 200MG BASE</u>	<u>A200363 004</u>	Mar 27, 2012
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A200363 005</u>	Mar 27, 2012
<u>AB</u>		<u>EQ 400MG BASE</u>	<u>A200363 006</u>	Mar 27, 2012
<u>AB</u>	UNICHEM LABS LTD	<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>AB</u>	ZENNOVA	<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

SEROQUEL

<u>AB</u>	+!	ASTRAZENECA PHARMS	<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>	ANCHEN PHARMS	<u>EQ 150MG BASE</u>	<u>A090757 001</u>	Dec 01, 2017
<u>AB</u>		<u>EQ 200MG BASE</u>	<u>A090757 002</u>	Dec 01, 2017
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A090757 003</u>	Dec 01, 2017
<u>AB</u>		<u>EQ 400MG BASE</u>	<u>A090757 004</u>	Dec 01, 2017
<u>AB</u>	AUROBINDO PHARMA LTD	<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>	INTELLIPHARMACEUTICS	<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

PRESCRIPTION DRUG PRODUCT LIST

QUETIAPINE FUMARATE

TABLET, EXTENDED RELEASE;ORAL

QUETIAPINE FUMARATE

<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>	PAR PHARM	<u>EQ 50MG BASE</u>	<u>A090482 001</u>	May 09, 2017
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A090482 002</u>	May 09, 2017
<u>AB</u>		<u>EQ 200MG BASE</u>	<u>A090482 003</u>	May 09, 2017
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A090482 004</u>	May 09, 2017
<u>AB</u>		<u>EQ 400MG BASE</u>	<u>A090482 005</u>	May 09, 2017
<u>AB</u>	PHARMADAX INC	<u>EQ 50MG BASE</u>	<u>A206260 001</u>	May 09, 2017
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A206260 002</u>	May 09, 2017
<u>AB</u>		<u>EQ 200MG BASE</u>	<u>A206260 003</u>	May 09, 2017
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A206260 004</u>	May 09, 2017
<u>AB</u>		<u>EQ 400MG BASE</u>	<u>A206260 005</u>	May 09, 2017
<u>AB</u>	SCIEGEN PHARMS INC	<u>EQ 50MG BASE</u>	<u>A209635 005</u>	Nov 16, 2018
<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

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

ACCUPRIL

<u>AB</u>	+	PFIZER PHARMS	<u>EQ 5MG BASE</u>	<u>N019885 001</u>	Nov 19, 1991
<u>AB</u>	+		<u>EQ 10MG BASE</u>	<u>N019885 002</u>	Nov 19, 1991
<u>AB</u>	+		<u>EQ 20MG BASE</u>	<u>N019885 003</u>	Nov 19, 1991
<u>AB</u>	+	!	<u>EQ 40MG BASE</u>	<u>N019885 004</u>	Nov 19, 1991

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>		INVAGEN PHARMS	<u>EQ 5MG BASE</u>	<u>A078457 001</u>	Aug 24, 2007
<u>AB</u>			<u>EQ 10MG BASE</u>	<u>A078457 002</u>	Aug 24, 2007
<u>AB</u>			<u>EQ 20MG BASE</u>	<u>A078457 003</u>	Aug 24, 2007
<u>AB</u>			<u>EQ 40MG BASE</u>	<u>A078457 004</u>	Aug 24, 2007
<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>		MYLAN	<u>EQ 5MG BASE</u>	<u>A076694 001</u>	Dec 23, 2004
<u>AB</u>			<u>EQ 10MG BASE</u>	<u>A076694 002</u>	Dec 23, 2004
<u>AB</u>			<u>EQ 20MG BASE</u>	<u>A076694 003</u>	Dec 23, 2004
<u>AB</u>			<u>EQ 40MG BASE</u>	<u>A076694 004</u>	Dec 23, 2004
<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

!	SUN PHARM INDUSTRIES	324MG	<u>A089338 001</u>	Feb 11, 1987
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QUINIDINE SULFATE

TABLET;ORAL

QUINIDINE SULFATE

<u>AB</u>		SANDOZ	<u>200MG</u>	<u>A088072 002</u>	
<u>AB</u>			<u>300MG</u>	<u>A088072 001</u>	Sep 26, 1983
<u>AB</u>	!	WATSON LABS	<u>200MG</u>	<u>A083288 001</u>	
<u>AB</u>	!		<u>300MG</u>	<u>A085583 001</u>	

PRESCRIPTION DRUG PRODUCT LIST

QUININE SULFATE

CAPSULE;ORAL

QUALAQUIN

AB	+ !	SUN PHARM INDUSTRIES	324MG	N021799 001	Aug 12, 2005
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QUININE SULFATE

AB		AMNEAL PHARMS	324MG	A203729 001	Jul 15, 2015
AB		LUPIN LTD	324MG	A203112 001	Apr 24, 2015
AB		NOVAST LABS	324MG	A204372 001	Jul 22, 2015
AB		TEVA PHARMS	324MG	A091661 001	Sep 28, 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

AB	+ !	EISAI INC	20MG	N020973 002	Aug 19, 1999
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RABEPRAZOLE SODIUM

AB		ALKEM LABS LTD	20MG	A208644 001	Apr 24, 2018
AB		AMNEAL PHARMS	20MG	A204179 001	Jul 31, 2015
AB		AUROBINDO PHARMA LTD	20MG	A205761 001	Feb 17, 2017
AB		DR REDDYS	20MG	A076824 001	Nov 08, 2013
AB		LANNETT CO INC	20MG	A090678 001	Nov 08, 2013
AB		LUPIN LTD	20MG	A078964 001	Nov 08, 2013
AB		RUBICON	20MG	A204237 001	Nov 18, 2015
AB		TEVA PHARMS USA	20MG	A076822 001	Nov 08, 2013
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
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RALOXIFENE HYDROCHLORIDE

AB		AMNEAL PHARMS	60MG	A208206 001	Apr 08, 2016
AB		AUROBINDO PHARMA LTD	60MG	A204310 001	Aug 28, 2015
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

+! MERCK SHARP DOHME

EQ 100MG BASE/PACKET

N205786 001 Dec 20, 2013

TABLET;ORAL

ISENTRESS

+! MERCK SHARP DOHME

EQ 400MG BASE

N022145 001 Oct 12, 2007

ISENTRESS HD

+! MERCK SHARP DOHME

EQ 600MG BASE

N022145 002 May 26, 2017

TABLET, CHEWABLE;ORAL

ISENTRESS

+ MERCK SHARP DOHME

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		DR REDDYS LABS SA	8MG	A091693 001	Jul 26, 2013
AB		ZYDUS PHARMS	8MG	A211567 001	Jul 22, 2019

ROZEREM

AB	+ !	TAKEDA PHARMS USA	8MG	N021782 001	Jul 22, 2005
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PRESCRIPTION DRUG PRODUCT LIST

3-376 (of 453)

RAMIPRIL

CAPSULE; ORAL

ALTACE

<u>AB</u>	+	KING PHARMS LLC	<u>1.25MG</u>	<u>N019901 001</u>	Jan 28, 1991
<u>AB</u>	+		<u>2.5MG</u>	<u>N019901 002</u>	Jan 28, 1991
<u>AB</u>	+		<u>5MG</u>	<u>N019901 003</u>	Jan 28, 1991
<u>AB</u>	+		<u>10MG</u>	<u>N019901 004</u>	Jan 28, 1991

RAMIPRIL

<u>AB</u>		ACCORD HLTHCARE	<u>1.25MG</u>	<u>A202392 001</u>	Apr 15, 2014
<u>AB</u>			<u>2.5MG</u>	<u>A202392 002</u>	Apr 15, 2014
<u>AB</u>			<u>5MG</u>	<u>A202392 003</u>	Apr 15, 2014
<u>AB</u>			<u>10MG</u>	<u>A202392 004</u>	Apr 15, 2014
<u>AB</u>		APOTEX	<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>		AUROBINDO PHARMA LTD	<u>1.25MG</u>	<u>A091604 001</u>	Jun 08, 2011
<u>AB</u>			<u>2.5MG</u>	<u>A091604 002</u>	Jun 08, 2011
<u>AB</u>			<u>5MG</u>	<u>A091604 003</u>	Jun 08, 2011
<u>AB</u>			<u>10MG</u>	<u>A091604 004</u>	Jun 08, 2011
<u>AB</u>		CHARTWELL MOLECULAR	<u>1.25MG</u>	<u>A078745 001</u>	Jun 18, 2008
<u>AB</u>			<u>2.5MG</u>	<u>A078745 002</u>	Jun 18, 2008
<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>		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>		TEVA PHARMS	<u>1.25MG</u>	<u>A077470 001</u>	Jun 18, 2008
<u>AB</u>			<u>2.5MG</u>	<u>A077470 002</u>	Jun 18, 2008
<u>AB</u>			<u>5MG</u>	<u>A077470 003</u>	Jun 18, 2008
<u>AB</u>			<u>10MG</u>	<u>A077470 004</u>	Jun 18, 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>		AJANTA PHARMA LTD	<u>EQ 150MG BASE</u>	<u>A209859 001</u>	Sep 27, 2018
<u>AB</u>			<u>EQ 300MG BASE</u>	<u>A209859 002</u>	Sep 27, 2018
<u>AB</u>		APPCO	<u>EQ 150MG BASE</u>	<u>A211893 001</u>	Apr 05, 2019
<u>AB</u>			<u>EQ 300MG BASE</u>	<u>A211893 002</u>	Apr 05, 2019
<u>AB</u>		AUROBINDO PHARMA LTD	<u>EQ 150MG BASE</u>	<u>A211058 001</u>	Jul 16, 2018
<u>AB</u>			<u>EQ 300MG BASE</u>	<u>A211058 002</u>	Jul 16, 2018
<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>		NOVITIUM PHARMA	<u>EQ 150MG BASE</u>	<u>A210681 001</u>	Nov 23, 2018
<u>AB</u>			<u>EQ 300MG BASE</u>	<u>A210681 002</u>	Nov 23, 2018
<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

INJECTABLE; INJECTION

RANITIDINE HYDROCHLORIDE

<u>AP</u>		MYLAN LABS LTD	<u>EQ 25MG BASE/ML</u>	<u>A079076 001</u>	Jun 09, 2016
<u>AP</u>		WEST-WARD PHARMS INT	<u>EQ 25MG BASE/ML</u>	<u>A074777 001</u>	Mar 02, 2005
<u>AP</u>			<u>EQ 25MG BASE/ML</u>	<u>A077458 001</u>	Feb 16, 2006
<u>AP</u>		ZYDUS PHARMS USA INC	<u>EQ 25MG BASE/ML</u>	<u>A091534 001</u>	Feb 22, 2013

PRESCRIPTION DRUG PRODUCT LIST

RANITIDINE HYDROCHLORIDE

INJECTABLE; INJECTION

ZANTAC

<u>AP</u>	<u>+</u> !	TELIGENT	<u>EQ 25MG BASE/ML</u>	<u>N019090 001</u>	Oct 19, 1984
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SYRUP; ORAL

RANITIDINE HYDROCHLORIDE

<u>AA</u>		AMNEAL PHARMS	<u>EQ 15MG BASE/ML</u>	<u>A078312 001</u>	Sep 02, 2008
<u>AA</u>		ANDA REPOSITORY	<u>EQ 15MG BASE/ML</u>	<u>A090054 001</u>	Nov 15, 2010
<u>AA</u>		AUROBINDO PHARMA LTD	<u>EQ 15MG BASE/ML</u>	<u>A090623 001</u>	Jul 28, 2010
<u>AA</u>		BRECKENRIDGE	<u>EQ 15MG BASE/ML</u>	<u>A078684 001</u>	Aug 27, 2009
<u>AA</u>		HI TECH PHARMA	<u>EQ 15MG BASE/ML</u>	<u>A091078 001</u>	Mar 22, 2011
<u>AA</u>		LANNETT CO INC	<u>EQ 15MG BASE/ML</u>	<u>A078890 001</u>	Jul 01, 2010
<u>AA</u>			<u>EQ 15MG BASE/ML</u>	<u>A091288 001</u>	Dec 09, 2010
<u>AA</u>		NOSTRUM LABS INC	<u>EQ 15MG BASE/ML</u>	<u>A091091 001</u>	Sep 20, 2011
<u>AA</u>	<u>!</u>	PHARM ASSOC	<u>EQ 15MG BASE/ML</u>	<u>A077405 001</u>	Sep 21, 2007
<u>AA</u>		TARO	<u>EQ 15MG BASE/ML</u>	<u>A077476 001</u>	Jun 13, 2011

TABLET; ORAL

RANITIDINE HYDROCHLORIDE

<u>AB</u>		ACIC PHARMS	<u>EQ 150MG BASE</u>	<u>A203694 001</u>	Nov 30, 2017
<u>AB</u>			<u>EQ 300MG BASE</u>	<u>A203694 002</u>	Nov 30, 2017
<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>		HERITAGE PHARMA	<u>EQ 150MG BASE</u>	<u>A075165 001</u>	Sep 30, 1998
<u>AB</u>			<u>EQ 300MG BASE</u>	<u>A075165 002</u>	Sep 30, 1998
<u>AB</u>		PAR PHARM	<u>EQ 150MG BASE</u>	<u>A075180 001</u>	Jan 28, 1999
<u>AB</u>			<u>EQ 300MG BASE</u>	<u>A075180 002</u>	Jan 28, 1999
<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>		STRIDES PHARMA	<u>EQ 150MG BASE</u>	<u>A205512 001</u>	Aug 22, 2016
<u>AB</u>	<u>!</u>		<u>EQ 300MG BASE</u>	<u>A205512 002</u>	Aug 22, 2016
<u>AB</u>		VKT PHARMA PVT LTD	<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
<u>AB</u>		WOCKHARDT LTD	<u>EQ 150MG BASE</u>	<u>A075208 001</u>	Dec 17, 1998
<u>AB</u>			<u>EQ 300MG BASE</u>	<u>A075208 002</u>	Dec 17, 1998

RANOLAZINE

TABLET, EXTENDED RELEASE; ORAL

RANEXA

<u>AB</u>	<u>+</u>	GILEAD	<u>500MG</u>	<u>N021526 002</u>	Jan 27, 2006
<u>AB</u>	<u>+</u> !		<u>1GM</u>	<u>N021526 001</u>	Feb 12, 2007

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>		ANI PHARMS INC	<u>500MG</u>	<u>A210482 001</u>	Oct 29, 2019
<u>AB</u>			<u>1GM</u>	<u>A210482 002</u>	Oct 29, 2019
<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>		CIPLA	<u>500MG</u>	<u>A211291 001</u>	May 28, 2019
<u>AB</u>			<u>1GM</u>	<u>A211291 002</u>	May 28, 2019
<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>		LUPIN LTD	<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>		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
<u>AB</u>			<u>1GM</u>	<u>A211745 002</u>	Feb 27, 2020
<u>AB</u>		SCIEGEN PHARMS INC	<u>500MG</u>	<u>A211829 001</u>	Jun 04, 2019
<u>AB</u>			<u>1GM</u>	<u>A211829 002</u>	Jun 04, 2019
<u>AB</u>		SUN PHARM	<u>500MG</u>	<u>A211707 001</u>	May 28, 2019
<u>AB</u>			<u>1GM</u>	<u>A211707 002</u>	May 28, 2019

PRESCRIPTION DRUG PRODUCT LIST

RASAGILINE MESYLATE

TABLET; ORAL

AZILECT

<u>AB</u>	+	TEVA	<u>EQ 0.5MG BASE</u>	<u>N021641 001</u>	May 16, 2006
<u>AB</u>	+	!	<u>EQ 1MG BASE</u>	<u>N021641 002</u>	May 16, 2006

RASAGILINE MESYLATE

<u>AB</u>		ALKEM LABS LTD	<u>EQ 0.5MG BASE</u>	<u>A201889 001</u>	Oct 30, 2017
<u>AB</u>			<u>EQ 1MG BASE</u>	<u>A201889 002</u>	Oct 30, 2017
<u>AB</u>		INDOCO REMEDIES	<u>EQ 0.5MG BASE</u>	<u>A206153 001</u>	Oct 04, 2019
<u>AB</u>			<u>EQ 1MG BASE</u>	<u>A206153 002</u>	Oct 04, 2019
<u>AB</u>		MICRO LABS	<u>EQ 0.5MG BASE</u>	<u>A207004 001</u>	Mar 29, 2019
<u>AB</u>			<u>EQ 1MG BASE</u>	<u>A207004 002</u>	Mar 29, 2019
<u>AB</u>		MYLAN	<u>EQ 0.5MG BASE</u>	<u>A201971 001</u>	May 15, 2017
<u>AB</u>			<u>EQ 1MG BASE</u>	<u>A201971 002</u>	May 15, 2017
<u>AB</u>		ORCHID HLTHCARE	<u>EQ 0.5MG BASE</u>	<u>A201970 001</u>	Mar 15, 2016
<u>AB</u>			<u>EQ 1MG BASE</u>	<u>A201970 002</u>	Mar 15, 2016
<u>AB</u>		SANDOZ INC	<u>EQ 0.5MG BASE</u>	<u>A201892 001</u>	Jul 27, 2018
<u>AB</u>			<u>EQ 1MG BASE</u>	<u>A201892 002</u>	Jul 27, 2018
<u>AB</u>		WATSON LABS INC	<u>EQ 0.5MG BASE</u>	<u>A201823 001</u>	Jul 01, 2013
<u>AB</u>			<u>EQ 1MG BASE</u>	<u>A201823 002</u>	Jul 01, 2013

REGADENOSON

SOLUTION; INTRAVENOUS

LEXISCAN

+	!	ASTELLAS	0.4MG/5ML (0.08MG/ML)	N022161 001	Apr 10, 2008
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REGORAFENIB

TABLET; ORAL

STIVARGA

+	!	BAYER HLTHCARE	40MG	N203085 001	Sep 27, 2012
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REMIFENTANIL HYDROCHLORIDE

INJECTABLE; INJECTION

REMIFENTANIL HYDROCHLORIDE

<u>AP</u>		FRESENIUS KABI USA	<u>EQ 1MG BASE/VIAL</u>	<u>A206223 001</u>	Jan 16, 2018
<u>AP</u>			<u>EQ 2MG BASE/VIAL</u>	<u>A206223 002</u>	Jan 16, 2018
<u>AP</u>			<u>EQ 5MG BASE/VIAL</u>	<u>A206223 003</u>	Jan 16, 2018
<u>ULTIVA</u>					
<u>AP</u>	+	MYLAN INSTITUTIONAL	<u>EQ 1MG BASE/VIAL</u>	<u>N020630 001</u>	Jul 12, 1996
<u>AP</u>	+		<u>EQ 2MG BASE/VIAL</u>	<u>N020630 002</u>	Jul 12, 1996
<u>AP</u>	+	!	<u>EQ 5MG BASE/VIAL</u>	<u>N020630 003</u>	Jul 12, 1996

REPAGLINIDE

TABLET; ORAL

REPAGLINIDE

<u>AB</u>		ACTAVIS TOTOWA	<u>0.5MG</u>	<u>A090008 001</u>	Jan 22, 2014
<u>AB</u>			<u>1MG</u>	<u>A090008 002</u>	Jan 22, 2014
<u>AB</u>			<u>2MG</u>	<u>A090008 003</u>	Jan 22, 2014
<u>AB</u>		AUROBINDO PHARMA LTD	<u>0.5MG</u>	<u>A203820 001</u>	Jan 22, 2014
<u>AB</u>			<u>1MG</u>	<u>A203820 002</u>	Jan 22, 2014
<u>AB</u>		!	<u>2MG</u>	<u>A203820 003</u>	Jan 22, 2014
<u>AB</u>		BOSCOGEN	<u>0.5MG</u>	<u>A091517 001</u>	Apr 24, 2015
<u>AB</u>			<u>1MG</u>	<u>A091517 002</u>	Apr 24, 2015
<u>AB</u>			<u>2MG</u>	<u>A091517 003</u>	Apr 24, 2015
<u>AB</u>		CASI PHARMS INC	<u>0.5MG</u>	<u>A078555 001</u>	Nov 22, 2013
<u>AB</u>			<u>1MG</u>	<u>A078555 002</u>	Jan 22, 2014
<u>AB</u>			<u>2MG</u>	<u>A078555 003</u>	Jan 22, 2014
<u>AB</u>		PADDOCK LLC	<u>0.5MG</u>	<u>A201189 001</u>	Jul 17, 2013
<u>AB</u>			<u>1MG</u>	<u>A201189 002</u>	Jan 22, 2014
<u>AB</u>			<u>2MG</u>	<u>A201189 003</u>	Jan 22, 2014
<u>AB</u>		SUN PHARM INDS INC	<u>1MG</u>	<u>A077571 002</u>	Jul 11, 2013
<u>AB</u>			<u>2MG</u>	<u>A077571 003</u>	Jul 11, 2013

RETAPAMULIN

OINTMENT; TOPICAL

ALTABAX

+	!	ALMIRALL	1%	N022055 001	Apr 12, 2007
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REVEFENACIN

SOLUTION; INHALATION

YUPELRI

+	!	MYLAN IRELAND LTD	175MCG/3ML	N210598 001	Nov 09, 2018
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PRESCRIPTION DRUG PRODUCT LIST

RIBAVIRIN

CAPSULE; ORAL

REBETOL

AB	+	! MERCK SHARP DOHME	200MG	N020903	002	Jul 25, 2001
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RIBAVIRIN

AB		AUROBINDO PHARMA	200MG	A079117	001	Sep 17, 2009
AB		TEVA	200MG	A076277	001	Oct 04, 2004
AB		ZYDUS PHARMS USA	200MG	A077224	001	Oct 28, 2005

FOR SOLUTION; INHALATION

RIBAVIRIN

AN		NAVINTA LLC	6GM/VIAL	A207366	001	Oct 06, 2016
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VIRAZOLE

AN	+	! VALEANT PHARM INTL	6GM/VIAL	N018859	001	Dec 31, 1985
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TABLET; ORAL

RIBAVIRIN

AB		AUROBINDO PHARMA	200MG	A079111	001	Sep 17, 2009
AB		SANDOZ	200MG	A077743	001	Oct 03, 2006
AB		ZYDUS PHARMS USA	200MG	A077094	001	Dec 05, 2005

RIBOCICLIB SUCCINATE

TABLET; ORAL

KISQALI

+	!	NOVARTIS	EQ 200MG BASE	N209092	001	Mar 13, 2017
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RIBOFLAVIN 5'-PHOSPHATE SODIUM

SOLUTION/DROPS; OPHTHALMIC

PHOTREXA

+	!	AVEDRO INC	0.146%	N203324	001	Apr 15, 2016
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PHOTREXA VISCOUS IN DEXTRAN 20%

+	!	AVEDRO INC	0.146%	N203324	002	Apr 15, 2016
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RIFABUTIN

CAPSULE; ORAL

MYCOBUTIN

AB	+	! PHARMACIA AND UPJOHN	150MG	N050689	001	Dec 23, 1992
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RIFABUTIN

AB		LUPIN LTD	150MG	A090033	001	Feb 24, 2014
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RIFAMPIN

CAPSULE; ORAL

RIFADIN

AB		SANOFI AVENTIS US	150MG	A062303	001	
AB	+	!	300MG	N050420	001	

RIFAMPIN

AB		AKORN	150MG	A065028	001	Mar 14, 2001
AB			300MG	A065028	002	Mar 14, 2001
AB		LANNETT CO INC	150MG	A065390	001	Mar 28, 2008
AB			300MG	A065390	002	Mar 28, 2008
AB		LUPIN PHARMS	150MG	A090034	001	Aug 21, 2013
AB			300MG	A090034	002	Aug 21, 2013
AB		SANDOZ	150MG	A064150	002	Jan 02, 1998
AB			300MG	A064150	001	May 28, 1997

RIMACTANE

AB		OXFORD PHARMS	300MG	N050429	001	
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INJECTABLE; INJECTION

RIFADIN

AP	+	!	600MG/VIAL	N050627	001	May 25, 1989
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RIFAMPIN

AP		AKORN	600MG/VIAL	A065502	001	Sep 21, 2010
AP		EMCURE PHARMS LTD	600MG/VIAL	A204101	001	Aug 18, 2014
AP		FRESENIUS KABI USA	600MG/VIAL	A091181	001	Aug 21, 2014
AP		HIKMA PHARMS	600MG/VIAL	A205039	001	Mar 03, 2016
AP		MYLAN LABS LTD	600MG/VIAL	A065421	001	May 22, 2008
AP		WATSON PHARMS TEVA	600MG/VIAL	A206736	001	Jan 19, 2016
AP		WEST-WARD PHARMS INT	600MG/VIAL	A064217	001	Oct 29, 1999

RIFAMYCIN SODIUM

TABLET, DELAYED RELEASE; ORAL

AEMCOLO

+	!	REDHILL	EQ 194MG BASE	N210910	001	Nov 16, 2018
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PRESCRIPTION DRUG PRODUCT LIST

RIFAPENTINE

TABLET;ORAL

PRIFTIN

+! SANOFI AVENTIS US 150MG N021024 001 Jun 22, 1998

RIFAXIMIN

TABLET;ORAL

XIFAXAN

+! SALIX PHARMS 200MG N021361 001 May 25, 2004

+! 550MG N022554 001 Mar 24, 2010

RILPIVIRINE HYDROCHLORIDE

TABLET;ORAL

EDURANT

+! JANSSEN PRODS EQ 25MG BASE N202022 001 May 20, 2011

RILUZOLE

FILM;ORAL

EXSERVAN

+! AQUESTIVE THERAP 50MG N212640 001 Nov 22, 2019

SUSPENSION;ORAL

TIGLUTIK KIT

+! ITALFARMACO SPA 50MG/10ML N209080 001 Sep 05, 2018

TABLET;ORAL

RILUTEK**AB** +! COVIS PHARMA BV **50MG** **N020599 001** Dec 12, 1995RILUZOLE**AB** ALKEM LABS LTD **50MG** **A204048 001** Mar 30, 2016**AB** APOTEX CORP **50MG** **A091300 001** Jun 18, 2013**AB** GLENMARK PHARMS LTD **50MG** **A091394 001** Jun 18, 2013**AB** IMPAX LABS **50MG** **A076173 001** Jan 29, 2003**AB** MYLAN PHARMS INC **50MG** **A203042 001** Jul 01, 2013**AB** STANDARD CHEM PHARM **50MG** **A206045 001** Dec 09, 2019**AB** SUN PHARM INDS LTD **50MG** **A091417 001** Jun 18, 2013RIMANTADINE HYDROCHLORIDE

TABLET;ORAL

FLUMADINE**AB** +! SUN PHARM INDS INC **100MG** **N019649 001** Sep 17, 1993RIMANTADINE HYDROCHLORIDE**AB** IMPAX LABS **100MG** **A076132 001** Aug 30, 2002RIMEGEPANT SULFATE

TABLET, ORALLY DISINTEGRATING;ORAL

NURTEC ODT

+! BIOHAVEN PHARM EQ 75MG BASE N212728 001 Feb 27, 2020

RIOCIGUAT

TABLET;ORAL

ADEMPAS

+ BAYER HLTHCARE 0.5MG N204819 001 Oct 08, 2013

+ 1MG N204819 002 Oct 08, 2013

+ 1.5MG N204819 003 Oct 08, 2013

+ 2MG N204819 004 Oct 08, 2013

+! 2.5MG N204819 005 Oct 08, 2013

RISEDRONATE SODIUM

TABLET;ORAL

ACTONEL**AB** + APIL **5MG** **N020835 002** Apr 14, 2000**AB** + **30MG** **N020835 001** Mar 27, 1998**AB** +! **35MG** **N020835 003** May 25, 2002**AB** +! **150MG** **N020835 005** Apr 22, 2008RISEDRONATE SODIUM**AB** APOTEX INC **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** **150MG** **A206768 001** Oct 21, 2016**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** ORCHID HLTHCARE **30MG** **A205280 001** May 13, 2019

PRESCRIPTION DRUG PRODUCT LIST

RISEDRONATE SODIUM

TABLET;ORAL

RISEDRONATE SODIUM

<u>AB</u>		<u>35MG</u>	<u>A205280 002</u>	May 13, 2019
<u>AB</u>	SUN PHARM	<u>5MG</u>	<u>A090886 001</u>	Nov 30, 2015
<u>AB</u>		<u>30MG</u>	<u>A090886 002</u>	Nov 30, 2015
<u>AB</u>		<u>35MG</u>	<u>A090886 003</u>	Nov 30, 2015
<u>AB</u>		<u>75MG</u>	<u>A090886 004</u>	Jun 10, 2014
<u>AB</u>		<u>150MG</u>	<u>A090886 005</u>	Jun 10, 2014
<u>AB</u>	TEVA PHARMS USA	<u>5MG</u>	<u>A077132 001</u>	Oct 05, 2007
<u>AB</u>		<u>30MG</u>	<u>A077132 002</u>	Oct 05, 2007
<u>AB</u>		<u>35MG</u>	<u>A077132 003</u>	Oct 05, 2007
<u>AB</u>		<u>150MG</u>	<u>A079215 001</u>	Jun 13, 2014

TABLET, DELAYED RELEASE;ORAL

ATELVIA

<u>AB</u>	+!	APIL	<u>35MG</u>	<u>N022560 001</u>	Oct 08, 2010
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RISEDRONATE SODIUM

<u>AB</u>	SUN PHARM	<u>35MG</u>	<u>A203925 001</u>	Jul 09, 2019
<u>AB</u>	TEVA PHARMS USA	<u>35MG</u>	<u>A203217 001</u>	May 18, 2015

RISPERIDONE

FOR SUSPENSION, EXTENDED RELEASE;SUBCUTANEOUS

PERSERIS KIT

+	INDIVIOR INC	90MG	N210655 001	Jul 27, 2018
+	!	120MG	N210655 002	Jul 27, 2018

INJECTABLE;INTRAMUSCULAR

RISPERDAL CONSTA

+	JANSSEN PHARMS	12.5MG/VIAL	N021346 004	Apr 12, 2007
+	!	25MG/VIAL	N021346 001	Oct 29, 2003
+		37.5MG/VIAL	N021346 002	Oct 29, 2003
+		50MG/VIAL	N021346 003	Oct 29, 2003

SOLUTION;ORAL

RISPERDAL

<u>AA</u>	+!	JANSSEN PHARMS	<u>1MG/ML</u>	<u>N020588 001</u>	Jun 10, 1996
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RISPERIDONE

<u>AA</u>		AMNEAL PHARMS	<u>1MG/ML</u>	<u>A091384 001</u>	May 25, 2011
<u>AA</u>		AUROBINDO PHARMA LTD	<u>1MG/ML</u>	<u>A078452 001</u>	Sep 04, 2009
<u>AA</u>		HIKMA	<u>1MG/ML</u>	<u>A076904 001</u>	Jul 29, 2009
<u>AA</u>		LANNETT CO INC	<u>1MG/ML</u>	<u>A079158 001</u>	Dec 03, 2010
<u>AA</u>		PHARM ASSOC	<u>1MG/ML</u>	<u>A077719 001</u>	Jul 29, 2009
<u>AA</u>		TARO	<u>1MG/ML</u>	<u>A090347 001</u>	Feb 07, 2011
<u>AA</u>		TRIS PHARMA INC	<u>1MG/ML</u>	<u>A079059 001</u>	Dec 12, 2012

TABLET;ORAL

RISPERDAL

<u>AB</u>	+	JANSSEN PHARMS	<u>0.25MG</u>	<u>N020272 008</u>	May 10, 1999
<u>AB</u>	+		<u>0.5MG</u>	<u>N020272 007</u>	Jan 27, 1999
<u>AB</u>	+!		<u>1MG</u>	<u>N020272 001</u>	Dec 29, 1993
<u>AB</u>	+		<u>2MG</u>	<u>N020272 002</u>	Dec 29, 1993
<u>AB</u>	+		<u>3MG</u>	<u>N020272 003</u>	Dec 29, 1993
<u>AB</u>	+		<u>4MG</u>	<u>N020272 004</u>	Dec 29, 1993

RISPERIDONE

<u>AB</u>		AJANTA PHARMA LTD	<u>0.25MG</u>	<u>A201003 001</u>	Aug 24, 2011
<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>		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>		CELLTRION	<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>		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

PRESCRIPTION DRUG PRODUCT LIST

RISPERIDONE

TABLET; ORAL

RISPERIDONE

<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>	MYLAN	<u>0.25MG</u>	<u>A076288 001</u>	Sep 15, 2008
<u>AB</u>		<u>0.5MG</u>	<u>A076288 002</u>	Sep 15, 2008
<u>AB</u>		<u>1MG</u>	<u>A076288 003</u>	Sep 15, 2008
<u>AB</u>		<u>2MG</u>	<u>A076288 004</u>	Sep 15, 2008
<u>AB</u>		<u>3MG</u>	<u>A076288 005</u>	Sep 15, 2008
<u>AB</u>		<u>4MG</u>	<u>A076288 006</u>	Sep 15, 2008
<u>AB</u>	OXFORD PHARMS	<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>	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
<u>AB</u>		<u>3MG</u>	<u>A079088 005</u>	Oct 30, 2008
<u>AB</u>		<u>4MG</u>	<u>A079088 006</u>	Oct 30, 2008
<u>AB</u>	ZYDUS PHARMS USA INC	<u>0.25MG</u>	<u>A078040 001</u>	Oct 16, 2008
<u>AB</u>		<u>0.5MG</u>	<u>A078040 002</u>	Oct 16, 2008
<u>AB</u>		<u>1MG</u>	<u>A078040 003</u>	Oct 16, 2008
<u>AB</u>		<u>2MG</u>	<u>A078040 004</u>	Oct 16, 2008
<u>AB</u>		<u>3MG</u>	<u>A078040 005</u>	Oct 16, 2008
<u>AB</u>		<u>4MG</u>	<u>A078040 006</u>	Oct 16, 2008

TABLET, ORALLY DISINTEGRATING; ORAL

RISPERDAL

<u>AB</u>	+	JANSSEN PHARMS	<u>0.5MG</u>	<u>N021444 001</u>	Apr 02, 2003
<u>AB</u>	+		<u>1MG</u>	<u>N021444 002</u>	Apr 02, 2003
<u>AB</u>	+		<u>2MG</u>	<u>N021444 003</u>	Apr 02, 2003
<u>AB</u>	+		<u>3MG</u>	<u>N021444 004</u>	Dec 23, 2004
<u>AB</u>	+		<u>4MG</u>	<u>N021444 005</u>	Dec 23, 2004

RISPERIDONE

<u>AB</u>	ACTAVIS LABS FL INC	<u>0.5MG</u>	<u>A076996 001</u>	Apr 19, 2011
<u>AB</u>		<u>1MG</u>	<u>A076996 002</u>	Apr 19, 2011
<u>AB</u>		<u>2MG</u>	<u>A076996 003</u>	Apr 19, 2011
<u>AB</u>		<u>3MG</u>	<u>A076996 004</u>	Apr 19, 2011

PRESCRIPTION DRUG PRODUCT LIST

RISPERIDONE

TABLET, ORALLY DISINTEGRATING;ORAL

RISPERIDONE

AB		<u>4MG</u>	<u>A076996</u>	<u>005</u>	Apr 19, 2011
AB	DR REDDYS LABS LTD	<u>0.5MG</u>	<u>A077328</u>	<u>001</u>	Feb 24, 2009
AB		<u>1MG</u>	<u>A077328</u>	<u>002</u>	Oct 05, 2009
AB		<u>2MG</u>	<u>A077328</u>	<u>003</u>	Feb 24, 2009
AB		<u>3MG</u>	<u>A077328</u>	<u>004</u>	Nov 30, 2009
AB		<u>4MG</u>	<u>A077328</u>	<u>005</u>	Nov 30, 2009
AB	HERITAGE PHARMA	<u>0.5MG</u>	<u>A076908</u>	<u>001</u>	Mar 12, 2012
AB		<u>1MG</u>	<u>A076908</u>	<u>002</u>	Mar 12, 2012
AB		<u>2MG</u>	<u>A076908</u>	<u>003</u>	Mar 12, 2012
AB	JUBILANT GENERICS	<u>0.5MG</u>	<u>A090839</u>	<u>001</u>	Nov 04, 2011
AB		<u>1MG</u>	<u>A090839</u>	<u>002</u>	Nov 04, 2011
AB		<u>2MG</u>	<u>A090839</u>	<u>003</u>	Nov 04, 2011
AB		<u>3MG</u>	<u>A090839</u>	<u>004</u>	Nov 04, 2011
AB		<u>4MG</u>	<u>A090839</u>	<u>005</u>	Nov 04, 2011
AB	PAR PHARM	<u>0.5MG</u>	<u>A077494</u>	<u>002</u>	Apr 30, 2009
AB		<u>1MG</u>	<u>A077494</u>	<u>003</u>	Oct 26, 2009
AB		<u>2MG</u>	<u>A077494</u>	<u>004</u>	Apr 30, 2009
AB		<u>3MG</u>	<u>A077494</u>	<u>005</u>	Apr 30, 2009
AB		<u>4MG</u>	<u>A077494</u>	<u>006</u>	Apr 30, 2009
AB	SANDOZ	<u>0.5MG</u>	<u>A078116</u>	<u>001</u>	Dec 22, 2009
AB		<u>1MG</u>	<u>A078116</u>	<u>002</u>	Dec 22, 2009
AB		<u>2MG</u>	<u>A078116</u>	<u>003</u>	Dec 22, 2009
AB		<u>3MG</u>	<u>A078116</u>	<u>004</u>	Dec 22, 2009
AB		<u>4MG</u>	<u>A078116</u>	<u>005</u>	Dec 22, 2009
AB	SUN PHARM INDS LTD	<u>0.5MG</u>	<u>A077542</u>	<u>001</u>	Aug 06, 2010
AB		<u>0.5MG</u>	<u>A078464</u>	<u>001</u>	Apr 08, 2013
AB		<u>1MG</u>	<u>A077542</u>	<u>002</u>	Aug 06, 2010
AB		<u>1MG</u>	<u>A078464</u>	<u>002</u>	Apr 08, 2013
AB		<u>2MG</u>	<u>A077542</u>	<u>003</u>	Aug 06, 2010
AB		<u>2MG</u>	<u>A078464</u>	<u>003</u>	Apr 08, 2013
AB		<u>3MG</u>	<u>A078464</u>	<u>004</u>	Apr 08, 2013
AB		<u>3MG</u>	<u>A078474</u>	<u>001</u>	Aug 06, 2010
AB		<u>4MG</u>	<u>A078464</u>	<u>005</u>	Apr 08, 2013
AB		<u>4MG</u>	<u>A078474</u>	<u>002</u>	Aug 06, 2010
AB	ZYDUS PHARMS USA	<u>0.5MG</u>	<u>A078516</u>	<u>001</u>	May 01, 2009
AB		<u>2MG</u>	<u>A078516</u>	<u>003</u>	May 01, 2009
	PAR PHARM	0.25MG	A077494	001	Apr 30, 2009

RITONAVIR

POWDER;ORAL

NORVIR

+! ABBVIE INC

100MG/PACKET

N209512 001 Jun 07, 2017

SOLUTION;ORAL

NORVIR

+! ABBVIE

80MG/ML

N020659 001 Mar 01, 1996

TABLET;ORAL

NORVIR

AB +! ABBVIE **100MG** **N022417** **001** Feb 10, 2010

RITONAVIR

AB AMNEAL PHARMS LLC **100MG** **A208890** **001** Sep 17, 2018

AB AUROBINDO PHARMA **100MG** **A206614** **001** Sep 17, 2018

AB LTD HETERO LABS LTD III **100MG** **A204587** **001** Sep 17, 2018

AB HIKMA **100MG** **A202573** **001** Jan 15, 2015

RIVAROXABAN

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

EXELON

AB + NOVARTIS **4.6MG/24HR** **N022083** **001** Jul 06, 2007

AB +! **9.5MG/24HR** **N022083** **002** Jul 06, 2007

AB + **13.3MG/24HR** **N022083** **005** Aug 31, 2012

RIVASTIGMINE

AB ALVOGEN **4.6MG/24HR** **A204403** **001** Sep 03, 2015

PRESCRIPTION DRUG PRODUCT LIST

3-384 (of 453)

RIVASTIGMINE

FILM, EXTENDED RELEASE;TRANSDERMAL

RIVASTIGMINE

<u>AB</u>		<u>9.5MG/24HR</u>	<u>A204403 002</u>	Sep 03, 2015
<u>AB</u>		<u>13.3MG/24HR</u>	<u>A204403 003</u>	Aug 31, 2015
<u>AB</u>	AMNEAL PHARMS	<u>4.6MG/24HR</u>	<u>A207308 001</u>	Jan 08, 2019
<u>AB</u>		<u>9.5MG/24HR</u>	<u>A207308 002</u>	Jan 08, 2019
<u>AB</u>		<u>13.3MG/24HR</u>	<u>A207308 003</u>	Jan 08, 2019
<u>AB</u>	BRECKENRIDGE	<u>4.6MG/24HR</u>	<u>A209063 001</u>	Nov 26, 2019
<u>AB</u>		<u>9.5MG/24HR</u>	<u>A209063 002</u>	Nov 26, 2019
<u>AB</u>		<u>13.3MG/24HR</u>	<u>A209063 003</u>	Nov 26, 2019
<u>AB</u>	MYLAN TECHNOLOGIES	<u>4.6MG/24HR</u>	<u>A205622 001</u>	Jun 20, 2018
<u>AB</u>		<u>9.5MG/24HR</u>	<u>A205622 002</u>	Jun 20, 2018
<u>AB</u>		<u>13.3MG/24HR</u>	<u>A205622 003</u>	Jun 20, 2018
<u>AB</u>	ZYDUS NOVELTECH INC	<u>4.6MG/24HR</u>	<u>A206318 001</u>	Mar 04, 2019
<u>AB</u>		<u>9.5MG/24HR</u>	<u>A206318 002</u>	Mar 04, 2019
<u>AB</u>		<u>13.3MG/24HR</u>	<u>A206318 003</u>	Mar 04, 2019

RIVASTIGMINE TARTRATE

CAPSULE;ORAL

RIVASTIGMINE TARTRATE

<u>AB</u>	ALEMBIC PHARMS LTD	<u>EQ 1.5MG BASE</u>	<u>A091689 001</u>	Jun 12, 2012
<u>AB</u>		<u>EQ 3MG BASE</u>	<u>A091689 002</u>	Jun 12, 2012
<u>AB</u>		<u>EQ 4.5MG BASE</u>	<u>A091689 003</u>	Jun 12, 2012
<u>AB</u>		<u>EQ 6MG BASE</u>	<u>A091689 004</u>	Jun 12, 2012
<u>AB</u>	APOTEX INC	<u>EQ 1.5MG BASE</u>	<u>A091072 001</u>	May 16, 2013
<u>AB</u>		<u>EQ 3MG BASE</u>	<u>A091072 002</u>	May 16, 2013
<u>AB</u>		<u>EQ 4.5MG BASE</u>	<u>A091072 003</u>	May 16, 2013
<u>AB</u>		<u>EQ 6MG BASE</u>	<u>A091072 004</u>	May 16, 2013
<u>AB</u>	AUROBINDO PHARMA LTD	<u>EQ 1.5MG BASE</u>	<u>A204572 001</u>	Mar 25, 2016
<u>AB</u>		<u>EQ 3MG BASE</u>	<u>A204572 002</u>	Mar 25, 2016
<u>AB</u>		<u>EQ 4.5MG BASE</u>	<u>A204572 003</u>	Mar 25, 2016
<u>AB</u>		<u>EQ 6MG BASE</u>	<u>A204572 004</u>	Mar 25, 2016
<u>AB</u>	CADILA PHARMS LTD	<u>EQ 1.5MG BASE</u>	<u>A203844 001</u>	Feb 13, 2017
<u>AB</u>		<u>EQ 3MG BASE</u>	<u>A203844 002</u>	Feb 13, 2017
<u>AB</u>		<u>EQ 4.5MG BASE</u>	<u>A203844 003</u>	Feb 13, 2017
<u>AB</u>		<u>EQ 6MG BASE</u>	<u>A203844 004</u>	Feb 13, 2017
<u>AB</u>	CHARTWELL RX	<u>EQ 1.5MG BASE</u>	<u>A207797 001</u>	Sep 28, 2017
<u>AB</u>		<u>EQ 3MG BASE</u>	<u>A207797 002</u>	Sep 28, 2017
<u>AB</u>		<u>EQ 4.5MG BASE</u>	<u>A207797 003</u>	Sep 28, 2017
<u>AB</u>		<u>EQ 6MG BASE</u>	<u>A207797 004</u>	Sep 28, 2017
<u>AB</u>	! DR REDDYS LABS INC	<u>EQ 1.5MG BASE</u>	<u>A077130 001</u>	Oct 31, 2007
<u>AB</u>		<u>EQ 3MG BASE</u>	<u>A077130 002</u>	Oct 31, 2007
<u>AB</u>		<u>EQ 4.5MG BASE</u>	<u>A077130 003</u>	Oct 31, 2007
<u>AB</u>	!	<u>EQ 6MG BASE</u>	<u>A077130 004</u>	Oct 31, 2007
<u>AB</u>	MACLEODS PHARMS LTD	<u>EQ 1.5MG BASE</u>	<u>A203148 001</u>	Aug 22, 2014
<u>AB</u>		<u>EQ 3MG BASE</u>	<u>A203148 002</u>	Aug 22, 2014
<u>AB</u>		<u>EQ 4.5MG BASE</u>	<u>A203148 003</u>	Aug 22, 2014
<u>AB</u>		<u>EQ 6MG BASE</u>	<u>A203148 004</u>	Aug 22, 2014
<u>AB</u>	ORCHID HLTHCARE	<u>EQ 1.5MG BASE</u>	<u>A090879 001</u>	Jun 10, 2015
<u>AB</u>		<u>EQ 3MG BASE</u>	<u>A090879 002</u>	Jun 10, 2015
<u>AB</u>		<u>EQ 4.5MG BASE</u>	<u>A090879 003</u>	Jun 10, 2015
<u>AB</u>		<u>EQ 6MG BASE</u>	<u>A090879 004</u>	Jun 10, 2015
<u>AB</u>	SUN PHARM	<u>EQ 1.5MG BASE</u>	<u>A077131 001</u>	Oct 22, 2007
<u>AB</u>		<u>EQ 3MG BASE</u>	<u>A077131 002</u>	Oct 22, 2007
<u>AB</u>		<u>EQ 4.5MG BASE</u>	<u>A077131 003</u>	Oct 22, 2007
<u>AB</u>		<u>EQ 6MG BASE</u>	<u>A077131 004</u>	Oct 22, 2007
<u>AB</u>	WATSON LABS	<u>EQ 1.5MG BASE</u>	<u>A077129 001</u>	Jan 08, 2008
<u>AB</u>		<u>EQ 3MG BASE</u>	<u>A077129 002</u>	Jan 08, 2008
<u>AB</u>		<u>EQ 4.5MG BASE</u>	<u>A077129 003</u>	Jan 08, 2008
<u>AB</u>		<u>EQ 6MG BASE</u>	<u>A077129 004</u>	Jan 08, 2008

RIZATRIPTAN BENZOATE

TABLET;ORAL

MAXALT

<u>AB</u>	+! MERCK	<u>EQ 10MG BASE</u>	<u>N020864 002</u>	Jun 29, 1998
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RIZATRIPTAN BENZOATE

<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>	CELLTRION	<u>EQ 5MG BASE</u>	<u>A077526 001</u>	Mar 26, 2013

PRESCRIPTION DRUG PRODUCT LIST

RIZATRIPTAN BENZOATE

TABLET; ORAL

RIZATRIPTAN BENZOATE

<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A077526 002</u>	Mar 26, 2013
<u>AB</u>	ECI PHARMS LLC	<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>	EMCURE PHARMS LTD	<u>EQ 5MG BASE</u>	<u>A204090 001</u>	Nov 26, 2013
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A204090 002</u>	Nov 26, 2013
<u>AB</u>	GLENMARK GENERICS	<u>EQ 5MG BASE</u>	<u>A201967 001</u>	Dec 31, 2012
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A201967 002</u>	Dec 31, 2012
<u>AB</u>	INVAGEN PHARMS	<u>EQ 5MG BASE</u>	<u>A204339 001</u>	Jul 01, 2013
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A204339 002</u>	Jul 01, 2013
<u>AB</u>	JUBILANT GENERICS	<u>EQ 5MG BASE</u>	<u>A203252 001</u>	Dec 31, 2014
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A203252 002</u>	Dec 31, 2014
<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>	NATCO PHARMA LTD	<u>EQ 5MG BASE</u>	<u>A200482 001</u>	Dec 31, 2012
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A200482 002</u>	Dec 31, 2012
<u>AB</u>	SANDOZ	<u>EQ 5MG BASE</u>	<u>A079230 001</u>	Dec 31, 2012
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A079230 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
<u>AB</u>	UNICHEM LABS LTD	<u>EQ 5MG BASE</u>	<u>A207836 001</u>	Mar 07, 2017
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A207836 002</u>	Mar 07, 2017

TABLET, ORALLY DISINTEGRATING; ORAL

MAXALT-MLT

<u>AB</u>	+! MERCK	<u>EQ 10MG BASE</u>	<u>N020865 002</u>	Jun 29, 1998
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RIZATRIPTAN BENZOATE

<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>	GLENMARK PHARMS LTD	<u>EQ 5MG BASE</u>	<u>A201914 001</u>	Jul 01, 2013
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A201914 002</u>	Jul 01, 2013
<u>AB</u>	JUBILANT GENERICS	<u>EQ 5MG BASE</u>	<u>A203334 001</u>	Oct 16, 2015
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A203334 002</u>	Oct 16, 2015
<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 BIOTEC LTD	<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
<u>AB</u>	SANDOZ	<u>EQ 5MG BASE</u>	<u>A078739 001</u>	Jul 01, 2013
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A078739 002</u>	Jul 01, 2013
<u>AB</u>	UNICHEM LABS LTD	<u>EQ 5MG BASE</u>	<u>A207835 001</u>	Mar 07, 2017
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A207835 002</u>	Mar 07, 2017

ROCURONIUM BROMIDE

INJECTABLE; INJECTION

ROCURONIUM BROMIDE

<u>AP</u>	AUROBINDO PHARMA LTD	<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>	GLAND PHARMA LTD	<u>50MG/5ML (10MG/ML)</u>	<u>A205656 001</u>	Apr 04, 2018
<u>AP</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>	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>	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 INC	<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>	SANOVEL ILAC	<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>	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>	TEVA PHARMS	<u>50MG/5ML (10MG/ML)</u>	<u>A078717 001</u>	Nov 26, 2008
<u>AP</u>		<u>100MG/10ML (10MG/ML)</u>	<u>A078717 002</u>	Nov 26, 2008
<u>AP</u>	WEST WARD PHARM	<u>50MG/5ML (10MG/ML)</u>	<u>A204679 001</u>	Feb 28, 2017

PRESCRIPTION DRUG PRODUCT LIST

ROCURONIUM BROMIDE

INJECTABLE; INJECTION

ROCURONIUM BROMIDE

CORP

AP 100MG/10ML (10MG/ML) **A204679 002** Feb 28, 2017

ROFLUMILAST

TABLET; ORAL

DALIRESP

AB **+**! ASTRAZENECA PHARMS 500MCG **N022522 001** Feb 28, 2011

ROFLUMILAST

AB BRECKENRIDGE 500MCG **A208236 001** Oct 03, 2018

AB HETERO LABS LTD III 500MCG **A208213 001** Nov 23, 2018

AB MICRO LABS 500MCG **A208180 001** Mar 22, 2019

AB TORRENT 500MCG **A208272 001** Aug 06, 2018

DALIRESP

+ ASTRAZENECA PHARMS 250MCG N022522 002 Jan 23, 2018

ROLAPITANT HYDROCHLORIDE

TABLET; ORAL

VARUBI

+! TERSERA THERAPS LLC EQ 90MG BASE N206500 001 Sep 01, 2015

ROMIDEPSIN

POWDER; INTRAVENOUS

ISTODAX

+! CELGENE 10MG/VIAL N022393 001 Nov 05, 2009

ROPINIROLE HYDROCHLORIDE

TABLET; ORAL

REQUIP

AB **+**! GLAXOSMITHKLINE LLC EQ 0.25MG BASE **N020658 001** Sep 19, 1997

AB **+** EQ 0.5MG BASE **N020658 002** Sep 19, 1997

AB **+** EQ 1MG BASE **N020658 003** Sep 19, 1997

AB **+** EQ 2MG BASE **N020658 004** Sep 19, 1997

AB **+** EQ 3MG BASE **N020658 006** Jan 27, 1999

AB **+** EQ 4MG BASE **N020658 007** Jan 27, 1999

AB **+** EQ 5MG BASE **N020658 005** Sep 19, 1997

ROPINIROLE HYDROCHLORIDE

AB ACCORD HLTHCARE EQ 0.25MG BASE **A204022 001** Feb 28, 2017

AB EQ 0.5MG BASE **A204022 002** Feb 28, 2017

AB EQ 1MG BASE **A204022 003** Feb 28, 2017

AB EQ 2MG BASE **A204022 004** Feb 28, 2017

AB EQ 3MG BASE **A204022 005** Feb 28, 2017

AB EQ 4MG BASE **A204022 006** Feb 28, 2017

AB EQ 5MG BASE **A204022 007** Feb 28, 2017

AB ALEMBIC LTD EQ 0.25MG BASE **A090429 001** Mar 24, 2010

AB EQ 0.5MG BASE **A090429 002** Mar 24, 2010

AB EQ 1MG BASE **A090429 003** Mar 24, 2010

AB EQ 2MG BASE **A090429 004** Mar 24, 2010

AB EQ 3MG BASE **A090429 005** Mar 24, 2010

AB EQ 4MG BASE **A090429 006** Mar 24, 2010

AB EQ 5MG BASE **A090429 007** Mar 24, 2010

AB CADILA EQ 0.25MG BASE **A090411 001** Jun 01, 2009

AB EQ 0.5MG BASE **A090411 002** Jun 01, 2009

AB EQ 1MG BASE **A090411 003** Jun 01, 2009

AB EQ 2MG BASE **A090411 004** Jun 01, 2009

AB EQ 3MG BASE **A090411 005** Jun 01, 2009

AB EQ 4MG BASE **A090411 006** Jun 01, 2009

AB EQ 5MG BASE **A090411 007** Jun 01, 2009

AB CELLTRION EQ 0.25MG BASE **A079050 001** May 29, 2008

AB EQ 0.5MG BASE **A079050 002** May 29, 2008

AB EQ 1MG BASE **A079050 003** May 29, 2008

AB EQ 2MG BASE **A079050 004** May 29, 2008

AB EQ 3MG BASE **A079050 005** May 29, 2008

AB EQ 4MG BASE **A079050 006** May 29, 2008

AB EQ 5MG BASE **A079050 007** May 29, 2008

AB GLENMARK GENERICS EQ 0.25MG BASE **A090135 001** Feb 25, 2010

AB EQ 0.5MG BASE **A090135 002** Feb 25, 2010

AB EQ 1MG BASE **A090135 003** Feb 25, 2010

AB EQ 2MG BASE **A090135 004** Feb 25, 2010

AB EQ 3MG BASE **A090135 005** Feb 25, 2010

AB EQ 4MG BASE **A090135 006** Feb 25, 2010

AB EQ 5MG BASE **A090135 007** Feb 25, 2010

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>	MYLAN	<u>EQ 0.25MG BASE</u>	<u>A078881 001</u>	May 05, 2008
<u>AB</u>		<u>EQ 0.5MG BASE</u>	<u>A078881 002</u>	May 05, 2008
<u>AB</u>		<u>EQ 1MG BASE</u>	<u>A078881 003</u>	May 05, 2008
<u>AB</u>		<u>EQ 2MG BASE</u>	<u>A078881 004</u>	May 05, 2008
<u>AB</u>		<u>EQ 3MG BASE</u>	<u>A078881 005</u>	May 05, 2008
<u>AB</u>		<u>EQ 4MG BASE</u>	<u>A078881 006</u>	May 05, 2008
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A078881 007</u>	May 19, 2008
<u>AB</u>	ORCHID HLTHCARE	<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

TABLET, EXTENDED RELEASE; ORAL

REQUIP XL

<u>AB</u>	+!	GLAXOSMITHKLINE LLC	<u>EQ 2MG BASE</u>	<u>N022008 001</u>	Jun 13, 2008
<u>AB</u>	+		<u>EQ 6MG BASE</u>	<u>N022008 006</u>	Apr 10, 2009
<u>AB</u>	+		<u>EQ 8MG BASE</u>	<u>N022008 004</u>	Jun 13, 2008
<u>AB</u>	+		<u>EQ 12MG BASE</u>	<u>N022008 005</u>	Oct 31, 2008

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 PHARMS LTD	<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>	CELLTRION	<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>	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
<u>AB</u>	WATSON LABS INC	<u>EQ 2MG BASE</u>	<u>A200431 001</u>	Jun 06, 2012
<u>AB</u>		<u>EQ 4MG BASE</u>	<u>A200431 002</u>	Jun 06, 2012
<u>AB</u>		<u>EQ 6MG BASE</u>	<u>A200431 003</u>	Jun 06, 2012
<u>AB</u>		<u>EQ 8MG BASE</u>	<u>A200431 004</u>	Jun 06, 2012
<u>AB</u>		<u>EQ 12MG BASE</u>	<u>A200431 005</u>	Jun 06, 2012

PRESCRIPTION DRUG PRODUCT LIST

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
<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>		AKORN	<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>		AKORN INC	<u>150MG/30ML (5MG/ML)</u>	<u>A203955 001</u>	Apr 11, 2016
<u>AP</u>		AUROBINDO PHARMA LTD	<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>		HOSPIRA	<u>20MG/10ML (2MG/ML)</u>	<u>A090194 001</u>	Sep 23, 2014
<u>AP</u>			<u>40MG/20ML (2MG/ML)</u>	<u>A090194 005</u>	Sep 23, 2014
<u>AP</u>			<u>100MG/10ML (10MG/ML)</u>	<u>A090194 004</u>	Sep 23, 2014
<u>AP</u>			<u>150MG/30ML (5MG/ML)</u>	<u>A090194 002</u>	Sep 23, 2014
<u>AP</u>			<u>150MG/20ML (7.5MG/ML)</u>	<u>A090194 003</u>	Sep 23, 2014
<u>AP</u>			<u>200MG/20ML (10MG/ML)</u>	<u>A090194 006</u>	Sep 23, 2014
<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>		MYLAN ASI	<u>40MG/20ML (2MG/ML)</u>	<u>A090318 001</u>	Sep 23, 2014
<u>AP</u>			<u>150MG/30ML (5MG/ML)</u>	<u>A090318 002</u>	Sep 23, 2014
<u>AP</u>			<u>150MG/20ML (7.5MG/ML)</u>	<u>A090318 003</u>	Sep 23, 2014
<u>AP</u>			<u>200MG/20ML (10MG/ML)</u>	<u>A090318 004</u>	Sep 23, 2014
<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>		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

ROSIGLITAZONE MALEATE

TABLET; ORAL

AVANDIA

+	SB PHARMCO	EQ 2MG BASE	N021071 002	May 25, 1999
+		EQ 4MG BASE	N021071 003	May 25, 1999

ROSUVASTATIN CALCIUM

CAPSULE; ORAL

EZALLOR

+	SUN PHARMA GLOBAL	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

CRESTOR

<u>AB</u>	+	IPR	<u>5MG</u>	<u>N021366 002</u>	Aug 12, 2003
<u>AB</u>	+		<u>10MG</u>	<u>N021366 003</u>	Aug 12, 2003
<u>AB</u>	+		<u>20MG</u>	<u>N021366 004</u>	Aug 12, 2003
<u>AB</u>	+		<u>40MG</u>	<u>N021366 005</u>	Aug 12, 2003

ROSUVASTATIN CALCIUM

<u>AB</u>		ACCORD HLTHCARE	<u>5MG</u>	<u>A206434 001</u>	Oct 31, 2016
<u>AB</u>			<u>10MG</u>	<u>A206434 002</u>	Oct 31, 2016
<u>AB</u>			<u>20MG</u>	<u>A206434 003</u>	Oct 31, 2016

PRESCRIPTION DRUG PRODUCT LIST

3-389 (of 453)

ROSUVASTATIN CALCIUM

TABLET; ORAL

ROSUVASTATIN CALCIUM

<u>AB</u>		<u>40MG</u>	<u>A206434</u>	<u>004</u>	Oct 31, 2016
<u>AB</u>	ALKEM LABS LTD	<u>5MG</u>	<u>A206465</u>	<u>001</u>	Mar 21, 2017
<u>AB</u>		<u>10MG</u>	<u>A206465</u>	<u>002</u>	Mar 21, 2017
<u>AB</u>		<u>20MG</u>	<u>A206465</u>	<u>003</u>	Mar 21, 2017
<u>AB</u>		<u>40MG</u>	<u>A206465</u>	<u>004</u>	Mar 21, 2017
<u>AB</u>	ALLIED	<u>5MG</u>	<u>A079168</u>	<u>001</u>	Jul 19, 2016
<u>AB</u>		<u>10MG</u>	<u>A079168</u>	<u>002</u>	Jul 19, 2016
<u>AB</u>		<u>20MG</u>	<u>A079168</u>	<u>003</u>	Jul 19, 2016
<u>AB</u>		<u>40MG</u>	<u>A079168</u>	<u>004</u>	Jul 19, 2016
<u>AB</u>	APOTEX INC	<u>5MG</u>	<u>A079145</u>	<u>001</u>	Jul 19, 2016
<u>AB</u>		<u>10MG</u>	<u>A079145</u>	<u>002</u>	Jul 19, 2016
<u>AB</u>		<u>20MG</u>	<u>A079145</u>	<u>003</u>	Jul 19, 2016
<u>AB</u>		<u>40MG</u>	<u>A079145</u>	<u>004</u>	Jul 19, 2016
<u>AB</u>	AUROBINDO PHARMA LTD	<u>5MG</u>	<u>A079170</u>	<u>001</u>	Jul 19, 2016
<u>AB</u>		<u>10MG</u>	<u>A079170</u>	<u>002</u>	Jul 19, 2016
<u>AB</u>		<u>20MG</u>	<u>A079170</u>	<u>003</u>	Jul 19, 2016
<u>AB</u>		<u>40MG</u>	<u>A079170</u>	<u>004</u>	Jul 19, 2016
<u>AB</u>	BIOCON LTD	<u>5MG</u>	<u>A207752</u>	<u>001</u>	Oct 31, 2016
<u>AB</u>		<u>10MG</u>	<u>A207752</u>	<u>002</u>	Oct 31, 2016
<u>AB</u>		<u>20MG</u>	<u>A207752</u>	<u>003</u>	Oct 31, 2016
<u>AB</u>		<u>40MG</u>	<u>A207752</u>	<u>004</u>	Oct 31, 2016
<u>AB</u>	CADILA PHARMS LTD	<u>5MG</u>	<u>A207453</u>	<u>001</u>	Nov 23, 2016
<u>AB</u>		<u>10MG</u>	<u>A207453</u>	<u>002</u>	Nov 23, 2016
<u>AB</u>		<u>20MG</u>	<u>A207453</u>	<u>003</u>	Nov 23, 2016
<u>AB</u>		<u>40MG</u>	<u>A207453</u>	<u>004</u>	Nov 23, 2016
<u>AB</u>	CHANGZHOU PHARM	<u>5MG</u>	<u>A207408</u>	<u>001</u>	Oct 31, 2016
<u>AB</u>		<u>10MG</u>	<u>A207408</u>	<u>002</u>	Oct 31, 2016
<u>AB</u>		<u>20MG</u>	<u>A207408</u>	<u>003</u>	Oct 31, 2016
<u>AB</u>		<u>40MG</u>	<u>A207408</u>	<u>004</u>	Oct 31, 2016
<u>AB</u>	GLENMARK PHARMS	<u>5MG</u>	<u>A079172</u>	<u>001</u>	Jul 19, 2016
<u>AB</u>		<u>10MG</u>	<u>A079172</u>	<u>002</u>	Jul 19, 2016
<u>AB</u>		<u>20MG</u>	<u>A079172</u>	<u>003</u>	Jul 19, 2016
<u>AB</u>		<u>40MG</u>	<u>A079172</u>	<u>004</u>	Jul 19, 2016
<u>AB</u>	HETERO LABS LTD V	<u>5MG</u>	<u>A207616</u>	<u>001</u>	Oct 31, 2016
<u>AB</u>		<u>10MG</u>	<u>A207616</u>	<u>002</u>	Oct 31, 2016
<u>AB</u>		<u>20MG</u>	<u>A207616</u>	<u>003</u>	Oct 31, 2016
<u>AB</u>		<u>40MG</u>	<u>A207616</u>	<u>004</u>	Oct 31, 2016
<u>AB</u>	JUBILANT GENERICS	<u>5MG</u>	<u>A207062</u>	<u>001</u>	Oct 31, 2016
<u>AB</u>		<u>10MG</u>	<u>A207062</u>	<u>002</u>	Oct 31, 2016
<u>AB</u>		<u>20MG</u>	<u>A207062</u>	<u>003</u>	Oct 31, 2016
<u>AB</u>		<u>40MG</u>	<u>A207062</u>	<u>004</u>	Oct 31, 2016
<u>AB</u>	LUPIN LTD	<u>5MG</u>	<u>A205587</u>	<u>001</u>	Jul 31, 2017
<u>AB</u>		<u>10MG</u>	<u>A205587</u>	<u>002</u>	Jul 31, 2017
<u>AB</u>		<u>20MG</u>	<u>A205587</u>	<u>003</u>	Jul 31, 2017
<u>AB</u>		<u>40MG</u>	<u>A205587</u>	<u>004</u>	Jul 31, 2017
<u>AB</u>	MSN	<u>5MG</u>	<u>A208898</u>	<u>001</u>	Nov 22, 2017
<u>AB</u>		<u>10MG</u>	<u>A208898</u>	<u>002</u>	Nov 22, 2017
<u>AB</u>		<u>20MG</u>	<u>A208898</u>	<u>003</u>	Nov 22, 2017
<u>AB</u>		<u>40MG</u>	<u>A208898</u>	<u>004</u>	Nov 22, 2017
<u>AB</u>	SANDOZ INC	<u>5MG</u>	<u>A079171</u>	<u>001</u>	Jul 19, 2016
<u>AB</u>		<u>10MG</u>	<u>A079171</u>	<u>002</u>	Jul 19, 2016
<u>AB</u>		<u>20MG</u>	<u>A079171</u>	<u>003</u>	Jul 19, 2016
<u>AB</u>		<u>40MG</u>	<u>A079171</u>	<u>004</u>	Jul 19, 2016
<u>AB</u>	SHANDONG	<u>5MG</u>	<u>A207375</u>	<u>001</u>	May 07, 2019
<u>AB</u>		<u>10MG</u>	<u>A207375</u>	<u>002</u>	May 07, 2019
<u>AB</u>		<u>20MG</u>	<u>A207375</u>	<u>003</u>	May 07, 2019
<u>AB</u>		<u>40MG</u>	<u>A207375</u>	<u>004</u>	May 07, 2019
<u>AB</u>	SUN PHARM	<u>5MG</u>	<u>A079169</u>	<u>001</u>	Jul 19, 2016
<u>AB</u>		<u>10MG</u>	<u>A079169</u>	<u>002</u>	Jul 19, 2016
<u>AB</u>		<u>20MG</u>	<u>A079169</u>	<u>003</u>	Jul 19, 2016
<u>AB</u>		<u>40MG</u>	<u>A079169</u>	<u>004</u>	Jul 19, 2016
<u>AB</u>	TORRENT	<u>5MG</u>	<u>A201619</u>	<u>001</u>	Oct 31, 2016
<u>AB</u>		<u>10MG</u>	<u>A201619</u>	<u>002</u>	Oct 31, 2016
<u>AB</u>		<u>20MG</u>	<u>A201619</u>	<u>003</u>	Oct 31, 2016
<u>AB</u>		<u>40MG</u>	<u>A201619</u>	<u>004</u>	Oct 31, 2016
<u>AB</u>	WATSON LABS INC	<u>5MG</u>	<u>A079167</u>	<u>001</u>	Apr 29, 2016
<u>AB</u>		<u>10MG</u>	<u>A079167</u>	<u>002</u>	Apr 29, 2016
<u>AB</u>		<u>20MG</u>	<u>A079167</u>	<u>003</u>	Apr 29, 2016
<u>AB</u>		<u>40MG</u>	<u>A079167</u>	<u>004</u>	Apr 29, 2016

PRESCRIPTION DRUG PRODUCT LISTROSUVASTATIN CALCIUM

TABLET; ORAL

ROSUVASTATIN CALCIUM

AB	ZHEJIANG YONGTAI	5MG	A212059 001	Nov 04, 2019
AB		10MG	A212059 002	Nov 04, 2019
AB		20MG	A212059 003	Nov 04, 2019
AB		40MG	A212059 004	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
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SOLUTION; INTRAVENOUS

RUBY-FILL

	JUBILANT DRAXIMAGE	N/A	N202153 001	Sep 30, 2016
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RUCAPARIB CAMSYLATE

TABLET; ORAL

RUBRACA

+	CLOVIS ONCOLOGY INC	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

AB	+	EISAI INC	40MG/ML	N201367 001	Mar 03, 2011
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RUFINAMIDE

AB		BIONPHARMA INC	40MG/ML	A211388 001	Apr 23, 2019
AB		HIKMA	40MG/ML	A207363 001	Apr 23, 2019

TABLET; ORAL

BANZEL

AB	+	EISAI INC	200MG	N021911 002	Nov 14, 2008
AB	+	!	400MG	N021911 003	Nov 14, 2008

RUFINAMIDE

AB		GLENMARK PHARMS LTD	200MG	A205075 001	May 16, 2016
AB			400MG	A205075 002	May 16, 2016
AB		HIKMA	200MG	A204988 001	May 16, 2016
AB			400MG	A204988 002	May 16, 2016

RUXOLITINIB PHOSPHATE

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
+		EQ 20MG BASE	N202192 004	Nov 16, 2011
+	!	EQ 25MG BASE	N202192 005	Nov 16, 2011

SACROSIDASE

SOLUTION; ORAL

SUCRAID

+	QOL MEDCL	8,500 IU/ML	N020772 001	Apr 09, 1998
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SACUBITRIL; VALSARTAN

TABLET; ORAL

ENTRESTO

+	NOVARTIS PHARMS CORP	24MG; 26MG	N207620 001	Jul 07, 2015
+		49MG; 51MG	N207620 002	Jul 07, 2015
+	!	97MG; 103MG	N207620 003	Jul 07, 2015

PRESCRIPTION DRUG PRODUCT LIST

SAFINAMIDE MESYLATE

TABLET; ORAL

XADAGO

+ US WORLDMEDS LLC

50MG

N207145 001 Mar 21, 2017

+!

100MG

N207145 002 Mar 21, 2017

SALMETEROL XINAFOATE

POWDER; INHALATION

SEREVENT

+! GLAXOSMITHKLINE

EQ 0.05MG BASE/INH

N020692 001 Sep 19, 1997

SAMARIUM SM-153 LEXIDRONAM PENTASODIUM

INJECTABLE; INJECTION

QUADRAMET

+! LANTHEUS MEDICAL

50mCi/ML

N020570 001 Mar 28, 1997

SAPROPTERIN DIHYDROCHLORIDE

POWDER; ORAL

KUVAN**AB +!** BIOMARIN PHARM **100MG/PACKET****N205065 001** Dec 19, 2013SAPROPTERIN DIHYDROCHLORIDE**AB** PAR PHARM INC **100MG/PACKET****A207207 001** Aug 20, 2019

KUVAN

+ BIOMARIN PHARM

500MG/PACKET

N205065 002 Oct 27, 2015

TABLET; ORAL

KUVAN

+! BIOMARIN PHARM

100MG

N022181 001 Dec 13, 2007

SAQUINAVIR MESYLATE

TABLET; ORAL

INVIRASE

+! HOFFMANN-LA ROCHE

EQ 500MG BASE

N021785 001 Dec 17, 2004

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

ONGLYZA

+ ASTRAZENECA AB

EQ 2.5MG BASE

N022350 001 Jul 31, 2009

+!

EQ 5MG BASE

N022350 002 Jul 31, 2009

SCOPOLAMINE

FILM, EXTENDED RELEASE; TRANSDERMAL

SCOPOLAMINE**AB** MYLAN TECHNOLOGIES **1MG/72HR****A203753 001** Jun 19, 2019**AB** PERRIGO PHARMS CO **1MG/72HR****A078830 001** Jan 30, 2015TRANSDERM SCOP**AB +!** GLAXOSMITHKLINE CON **1MG/72HR****N017874 001**SECNIDAZOLE

GRANULE; ORAL

SOLOSEC

+! LUPIN

2GM/PACKET

N209363 001 Sep 15, 2017

SECOBARBITAL SODIUM

CAPSULE; ORAL

SECONAL SODIUM

! VALEANT PHARMS

50MG

A086101 001 Oct 03, 1983

NORTH

!

100MG

A086101 002 Oct 03, 1983

SECRETIN SYNTHETIC HUMAN

FOR SOLUTION; INTRAVENOUS

CHIRHOSTIM

+! CHIRHOCLIN

16MCG/VIAL

N021256 001 Apr 09, 2004

+

40MCG/VIAL

N021256 002 Jun 21, 2007

PRESCRIPTION DRUG PRODUCT LIST

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		RISING	5MG	A206803 001	Apr 02, 2019

TABLET;ORAL

SELEGILINE HYDROCHLORIDE

AB	!	APOTEX INC	5MG	A074871 001	Jun 06, 1997
AB		BOSCOGEN	5MG	A074912 001	Apr 30, 1998

TABLET, ORALLY DISINTEGRATING;ORAL

ZELAPAR

+	!	BAUSCH	1.25MG	N021479 001	Jun 14, 2006
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SELENIOS ACID

SOLUTION;INTRAVENOUS

SELENIOS ACID

+	!	AM REGENT	EQ 600MCG BASE/10ML (EQ 60MCG BASE/ML)	N209379 001	Apr 30, 2019
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SELENIUM SULFIDE

LOTION;SHAMPOO;TOPICAL

SELENIUM SULFIDE

AT	!	PERRIGO NEW YORK	2.5%	A089996 001	Jan 10, 1991
AT		WOCKHARDT BIO AG	2.5%	A088228 001	Sep 01, 1983

SELEXIPAG

TABLET;ORAL

UPTRAIVI

+		ACTELION PHARMS LTD	0.2MG	N207947 001	Dec 21, 2015
+	!		0.4MG	N207947 002	Dec 21, 2015
+			0.6MG	N207947 003	Dec 21, 2015
+			0.8MG	N207947 004	Dec 21, 2015
+			1MG	N207947 005	Dec 21, 2015
+			1.2MG	N207947 006	Dec 21, 2015
+			1.4MG	N207947 007	Dec 21, 2015
+			1.6MG	N207947 008	Dec 21, 2015

SELINEXOR

TABLET;ORAL

XPOVIO

+	!	KARYOPHARM THERAPS	20MG	N212306 001	Jul 03, 2019
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SEMAGLUTIDE

SOLUTION;SUBCUTANEOUS

OZEMPIC

+	!	NOVO	2MG/1.5ML (1.34MG/ML)	N209637 001	Dec 05, 2017
+	!		4MG/3ML (1.34MG/ML)	N209637 002	Apr 09, 2019

TABLET;ORAL

RYBELSUS

+		NOVO	3MG	N213051 001	Sep 20, 2019
+			7MG	N213051 002	Sep 20, 2019
+	!		14MG	N213051 003	Sep 20, 2019

SERTACONAZOLE NITRATE

CREAM;TOPICAL

ERTACZO

+	!	BAUSCH	2%	N021385 001	Dec 10, 2003
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SERTRALINE HYDROCHLORIDE

CONCENTRATE;ORAL

SERTRALINE HYDROCHLORIDE

AA		AUROBINDO PHARMA	EQ 20MG BASE/ML	A078861 001	Oct 31, 2008
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ZOLOFT

AA	+	PFIZER	EQ 20MG BASE/ML	N020990 001	Dec 07, 1999
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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		APOTEX INC	EQ 25MG BASE	A076882 001	Feb 06, 2007

PRESCRIPTION DRUG PRODUCT LIST

SERTRALINE HYDROCHLORIDE

TABLET;ORAL

SERTRALINE HYDROCHLORIDE

<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A076882 002</u>	Feb 06, 2007
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A076882 003</u>	Feb 06, 2007
<u>AB</u>	AUROBINDO PHARMA	<u>EQ 25MG BASE</u>	<u>A077206 001</u>	Feb 06, 2007
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A077206 002</u>	Feb 06, 2007
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A077206 003</u>	Feb 06, 2007
<u>AB</u>	AUSTARPHARMA LLC	<u>EQ 25MG BASE</u>	<u>A078677 001</u>	Mar 04, 2009
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A078677 002</u>	Mar 04, 2009
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A078677 003</u>	Mar 04, 2009
<u>AB</u>	INVAGEN PHARMS	<u>EQ 25MG BASE</u>	<u>A077397 001</u>	Feb 06, 2007
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A077397 002</u>	Feb 06, 2007
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A077397 003</u>	Feb 06, 2007
<u>AB</u>	LUPIN	<u>EQ 25MG BASE</u>	<u>A077670 001</u>	Feb 06, 2007
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A077670 002</u>	Feb 06, 2007
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A077670 003</u>	Feb 06, 2007
<u>AB</u>	OXFORD PHARMS	<u>EQ 25MG BASE</u>	<u>A078175 001</u>	Jul 21, 2010
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A078175 002</u>	Jul 21, 2010
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A078175 003</u>	Jul 21, 2010
<u>AB</u>	SUN PHARM INDS LTD	<u>EQ 25MG BASE</u>	<u>A077977 001</u>	Feb 06, 2007
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A077977 002</u>	Feb 06, 2007
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A077977 003</u>	Feb 06, 2007
<u>AB</u>	TORRENT PHARMS	<u>EQ 25MG BASE</u>	<u>A077765 001</u>	Feb 06, 2007
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A077765 002</u>	Feb 06, 2007
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A077765 003</u>	Feb 06, 2007
<u>AB</u>	WOCKHARDT	<u>EQ 25MG BASE</u>	<u>A078403 001</u>	Jan 08, 2008
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A078403 002</u>	Jan 08, 2008
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A078403 003</u>	Jan 08, 2008

ZOLOFT

<u>AB</u>	+ PFIZER	<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
	SERTRALINE HYDROCHLORIDE			
	SUN PHARM INDS LTD	EQ 150MG BASE	A077977 004	Feb 06, 2007
		EQ 200MG BASE	A077977 005	Feb 06, 2007

SEVELAMER CARBONATE

FOR SUSPENSION;ORAL

RENVELA

<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
	<u>SEVELAMER CARBONATE</u>			
<u>AB</u>	AUROBINDO PHARMA LTD	<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>	DR REDDYS LABS LTD	<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

TABLET;ORAL

RENVELA

<u>AB</u>	+! SANOFI	<u>800MG</u>	<u>N022127 001</u>	Oct 19, 2007
	<u>SEVELAMER CARBONATE</u>			
<u>AB</u>	AMNEAL PHARMS CO	<u>800MG</u>	<u>A207288 001</u>	Nov 28, 2017
<u>AB</u>	AUROBINDO PHARMA LTD	<u>800MG</u>	<u>A207179 001</u>	Jul 17, 2017
<u>AB</u>	DR REDDYS LABS LTD	<u>800MG</u>	<u>A206094 001</u>	Sep 29, 2017
<u>AB</u>	IMPAX LABS INC	<u>800MG</u>	<u>A090975 001</u>	Oct 23, 2017
<u>AB</u>	INVAGEN PHARMS	<u>800MG</u>	<u>A203860 001</u>	Oct 26, 2017
<u>AB</u>	TWI PHARMS	<u>800MG</u>	<u>A200959 001</u>	Mar 20, 2018
<u>AB</u>	WILSHIRE PHARMS INC	<u>800MG</u>	<u>A204451 001</u>	Nov 29, 2018

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
	<u>SEVELAMER HYDROCHLORIDE</u>			
<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

PRESCRIPTION DRUG PRODUCT LIST

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>	SHANGHAI HENGRUI	<u>100%</u>	<u>A203793 001</u>	Nov 03, 2015

SOJOURN

<u>AN</u>	PIRAMAL CRITICAL	<u>100%</u>	<u>A077867 001</u>	May 02, 2007
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ULTANE

<u>AN</u>	+! ABBVIE	<u>100%</u>	<u>N020478 001</u>	Jun 07, 1995
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SILDENAFIL CITRATE

FOR SUSPENSION; ORAL

REVATIO

<u>AB</u>	+! PFIZER	<u>EQ 10MG BASE/ML</u>	<u>N203109 001</u>	Aug 30, 2012
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SILDENAFIL CITRATE

<u>AB</u>	AJANTA PHARMA LTD	<u>EQ 10MG BASE/ML</u>	<u>A212883 001</u>	Nov 27, 2019
<u>AB</u>	ALKEM LABS LTD	<u>EQ 10MG BASE/ML</u>	<u>A212440 001</u>	Nov 27, 2019
<u>AB</u>	AMNEAL PHARMS	<u>EQ 10MG BASE/ML</u>	<u>A211092 001</u>	Nov 27, 2019
<u>AB</u>	NOVITIUM PHARMA	<u>EQ 10MG BASE/ML</u>	<u>A212260 001</u>	May 31, 2019

SOLUTION; INTRAVENOUS

REVATIO

<u>AP</u>	+! PFIZER	<u>EQ 10MG BASE/12.5ML (EQ 0.8MG BASE/ML)</u>	<u>N022473 001</u>	Nov 18, 2009
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SILDENAFIL CITRATE

<u>AP</u>	AUROBINDO PHARMA LTD	<u>EQ 10MG BASE/12.5ML (EQ 0.8MG BASE/ML)</u>	<u>A203988 001</u>	Apr 01, 2015
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TABLET; ORAL

REVATIO

<u>AB</u>	+! PFIZER	<u>EQ 20MG BASE</u>	<u>N021845 001</u>	Jun 03, 2005
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SILDENAFIL CITRATE

<u>AB</u>	AJANTA PHARMA LTD	<u>EQ 20MG BASE</u>	<u>A210394 001</u>	May 04, 2018
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>A206401 001</u>	Oct 12, 2018
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A206401 002</u>	Oct 12, 2018
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A206401 003</u>	Oct 12, 2018
<u>AB</u>	AMNEAL PHARMS	<u>EQ 20MG BASE</u>	<u>A202025 001</u>	Feb 28, 2013
<u>AB</u>	AMNEAL PHARMS NY	<u>EQ 25MG BASE</u>	<u>A202023 001</u>	Jun 27, 2018
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A202023 002</u>	Jun 27, 2018
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A202023 003</u>	Jun 27, 2018
<u>AB</u>	APPCO	<u>25MG</u>	<u>A207178 001</u>	Mar 02, 2020
<u>AB</u>		<u>50MG</u>	<u>A207178 002</u>	Mar 02, 2020
<u>AB</u>		<u>100MG</u>	<u>A207178 003</u>	Mar 02, 2020
<u>AB</u>	AUROBINDO PHARMA LTD	<u>EQ 20MG BASE</u>	<u>A203963 001</u>	Nov 18, 2015
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>A203962 001</u>	Jun 11, 2018
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A203962 002</u>	Jun 11, 2018
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A203962 003</u>	Jun 11, 2018
<u>AB</u>	HEBEI CHANGSHAN	<u>EQ 20MG BASE</u>	<u>A202598 001</u>	Nov 06, 2012
<u>AB</u>	HETERO LABS LTD V	<u>EQ 20MG BASE</u>	<u>A203623 001</u>	Nov 26, 2014
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>A202659 001</u>	Jun 11, 2018
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A202659 002</u>	Jun 11, 2018
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A202659 003</u>	Jun 11, 2018
<u>AB</u>	LUPIN LTD	<u>EQ 25MG BASE</u>	<u>A212051 001</u>	Mar 22, 2019
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A212051 002</u>	Mar 22, 2019
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A212051 003</u>	Mar 22, 2019
<u>AB</u>	MACLEODS PHARMS LTD	<u>EQ 20MG BASE</u>	<u>A203814 001</u>	Dec 17, 2013
<u>AB</u>	MYLAN	<u>EQ 25MG BASE</u>	<u>A201171 001</u>	Mar 25, 2019
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A201171 002</u>	Mar 25, 2019
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A201171 003</u>	Mar 25, 2019
<u>AB</u>	MYLAN PHARMS INC	<u>EQ 20MG BASE</u>	<u>A201150 001</u>	Nov 09, 2012
<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>	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

PRESCRIPTION DRUG PRODUCT LIST

SILDENAFIL CITRATE

TABLET;ORAL

VIAGRA

AB	+	PFIZER INC	<u>EQ 25MG BASE</u>	<u>N020895 001</u>	Mar 27, 1998
AB	+		<u>EQ 50MG BASE</u>	<u>N020895 002</u>	Mar 27, 1998
AB	+		<u>EQ 100MG BASE</u>	<u>N020895 003</u>	Mar 27, 1998

SILODOSIN

CAPSULE;ORAL

RAPAFLO

AB	+	ALLERGAN	<u>4MG</u>	<u>N022206 001</u>	Oct 08, 2008
AB	+		<u>8MG</u>	<u>N022206 002</u>	Oct 08, 2008

SILODOSIN

AB		AJANTA PHARMA LTD	<u>4MG</u>	<u>A211060 001</u>	Dec 03, 2018
AB			<u>8MG</u>	<u>A211060 002</u>	Dec 03, 2018
AB		ALEMBIC PHARMS LTD	<u>4MG</u>	<u>A211731 001</u>	Nov 22, 2019
AB			<u>8MG</u>	<u>A211731 002</u>	Nov 22, 2019
AB		AMNEAL PHARMS CO	<u>4MG</u>	<u>A209745 001</u>	Dec 03, 2018
AB			<u>8MG</u>	<u>A209745 002</u>	Dec 03, 2018
AB		AUROBINDO PHARMA LTD	<u>4MG</u>	<u>A210626 001</u>	Dec 10, 2018
AB			<u>8MG</u>	<u>A210626 002</u>	Dec 10, 2018
AB		HETERO LABS LTD V	<u>4MG</u>	<u>A204793 001</u>	Feb 14, 2020
AB			<u>8MG</u>	<u>A204793 002</u>	Feb 14, 2020
AB		LUPIN LTD	<u>4MG</u>	<u>A206541 001</u>	Dec 03, 2018
AB			<u>8MG</u>	<u>A206541 002</u>	Dec 03, 2018
AB		MACLEODS PHARMS LTD	<u>4MG</u>	<u>A211166 001</u>	Dec 03, 2018
AB			<u>8MG</u>	<u>A211166 002</u>	Dec 03, 2018
AB		MSN	<u>4MG</u>	<u>A210687 001</u>	Dec 03, 2018
AB			<u>8MG</u>	<u>A210687 002</u>	Dec 03, 2018
AB		SANDOZ INC	<u>4MG</u>	<u>A204726 001</u>	Mar 31, 2017
AB			<u>8MG</u>	<u>A204726 002</u>	Mar 31, 2017

SILVER SULFADIAZINE

CREAM;TOPICAL

SILVADENE

AB	+	KING PHARMS LLC	<u>1%</u>	<u>N017381 001</u>	
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SSD

AB		DR REDDYS LA	<u>1%</u>	<u>N018578 001</u>	Feb 25, 1982
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THERMAZENE

AB		THEPHARMANETWORK LLC	<u>1%</u>	<u>N018810 001</u>	Dec 23, 1985
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SIMVASTATIN

SUSPENSION;ORAL

FLOLIPID

+	TCG FLUENT PHARMA	20MG/5ML	<u>N206679 001</u>	Apr 21, 2016
+		40MG/5ML	<u>N206679 002</u>	Apr 21, 2016

TABLET;ORAL

SIMVASTATIN

AB		ACCORD HLTHCARE	<u>5MG</u>	<u>A078155 005</u>	Apr 05, 2013
AB			<u>10MG</u>	<u>A078155 002</u>	Feb 26, 2008
AB			<u>20MG</u>	<u>A078155 003</u>	Feb 26, 2008
AB			<u>40MG</u>	<u>A078155 004</u>	Feb 26, 2008
AB			<u>80MG</u>	<u>A078155 001</u>	Feb 26, 2008
AB		AUROBINDO PHARMA	<u>5MG</u>	<u>A077691 001</u>	Dec 20, 2006
AB			<u>10MG</u>	<u>A077691 002</u>	Dec 20, 2006
AB			<u>20MG</u>	<u>A077691 003</u>	Dec 20, 2006
AB			<u>40MG</u>	<u>A077691 004</u>	Dec 20, 2006
AB			<u>80MG</u>	<u>A077691 005</u>	Dec 20, 2006
AB		BIOCON LIMITED	<u>5MG</u>	<u>A078034 001</u>	Dec 20, 2006
AB			<u>10MG</u>	<u>A078034 002</u>	Dec 20, 2006
AB			<u>20MG</u>	<u>A078034 003</u>	Dec 20, 2006
AB			<u>40MG</u>	<u>A078034 004</u>	Dec 20, 2006
AB			<u>80MG</u>	<u>A078034 005</u>	Dec 20, 2006
AB		DR REDDYS LABS INC	<u>5MG</u>	<u>A077752 005</u>	Jan 23, 2008
AB			<u>10MG</u>	<u>A077752 001</u>	Dec 20, 2006
AB			<u>20MG</u>	<u>A077752 002</u>	Dec 20, 2006
AB			<u>40MG</u>	<u>A077752 003</u>	Dec 20, 2006
AB			<u>80MG</u>	<u>A077752 004</u>	Dec 20, 2006
AB		HETERO LABS LTD III	<u>5MG</u>	<u>A200895 001</u>	Nov 25, 2014
AB			<u>10MG</u>	<u>A200895 002</u>	Nov 25, 2014
AB			<u>20MG</u>	<u>A200895 003</u>	Nov 25, 2014
AB			<u>40MG</u>	<u>A200895 004</u>	Nov 25, 2014

PRESCRIPTION DRUG PRODUCT LIST

SIMVASTATIN

TABLET; ORAL

SIMVASTATIN

<u>AB</u>		<u>80MG</u>	<u>A200895 005</u>	Nov 25, 2014
<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>	WATSON LABS TEVA	<u>5MG</u>	<u>A076685 001</u>	Dec 20, 2006
<u>AB</u>		<u>10MG</u>	<u>A076685 002</u>	Dec 20, 2006
<u>AB</u>		<u>20MG</u>	<u>A076685 003</u>	Dec 20, 2006
<u>AB</u>		<u>40MG</u>	<u>A076685 004</u>	Dec 20, 2006
<u>AB</u>		<u>80MG</u>	<u>A076685 005</u>	Dec 20, 2006
<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

ZOCOR

<u>AB</u>	+	MSD MERCK CO	<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
<u>AB</u>	+		<u>80MG</u>	<u>N019766 005</u>	Jul 10, 1998

SINCALIDE

INJECTABLE; INJECTION

KINEVAC

+! BRACCO

0.005MG/VIAL

N017697 001

SINECATECHINS

OINTMENT; TOPICAL

VEREGEN

+! FOUGERA PHARMS INC

15%

N021902 001 Oct 31, 2006

SIPONIMOD FUMARIC ACID

TABLET; ORAL

MAYZENT

+ NOVARTIS

EQ 0.25MG BASE

N209884 001 Mar 26, 2019

+!

EQ 2MG BASE

N209884 002 Mar 26, 2019

SIROLIMUS

SOLUTION; ORAL

RAPAMUNE

<u>AA</u>	+	PF PRISM CV	<u>1MG/ML</u>	<u>N021083 001</u>	Sep 15, 1999
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SIROLIMUS

<u>AA</u>		AMNEAL PHARMS LLC	<u>1MG/ML</u>	<u>A211212 001</u>	Oct 18, 2019
<u>AA</u>		APOTEX	<u>1MG/ML</u>	<u>A211406 001</u>	Oct 22, 2019
<u>AA</u>		NOVITIUM PHARMA	<u>1MG/ML</u>	<u>A211040 001</u>	Jan 28, 2019

TABLET; ORAL

RAPAMUNE

<u>AB</u>	+	PF PRISM CV	<u>0.5MG</u>	<u>N021110 004</u>	Jan 25, 2010
<u>AB</u>	+		<u>1MG</u>	<u>N021110 001</u>	Aug 25, 2000
<u>AB</u>	+		<u>2MG</u>	<u>N021110 002</u>	Aug 22, 2002

SIROLIMUS

<u>AB</u>		DR REDDYS LABS LTD	<u>1MG</u>	<u>A201578 001</u>	Oct 27, 2014
<u>AB</u>			<u>2MG</u>	<u>A201578 002</u>	Oct 27, 2014
<u>AB</u>		ZYDUS PHARMS	<u>0.5MG</u>	<u>A201676 003</u>	Jan 08, 2014

PRESCRIPTION DRUG PRODUCT LIST

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

INJECTABLE; INJECTION

SODIUM ACETATE

	FRESENIUS KABI USA	4MEQ/ML	A206687 001	Oct 30, 2017
+	!	HOSPIRA	N018893 001	May 04, 1983

SODIUM BENZOATE; SODIUM PHENYLACETATE

SOLUTION; INTRAVENOUS

AMMONUL

<u>AP</u>	+	MEDICIS	<u>10%;10% (5GM/50ML;5GM/50ML)</u>	<u>N020645 001</u>	Feb 17, 2005
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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>		NAVINTA LLC	<u>10%;10% (5GM/50ML;5GM/50ML)</u>	<u>A205880 001</u>	Aug 04, 2016

SODIUM BICARBONATE

INJECTABLE; INJECTION

SODIUM BICARBONATE

<u>AP</u>		EXELA PHARMA SCS LLC	<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>0.9MEQ/ML</u>	<u>A077394 001</u>	Nov 09, 2005
<u>AP</u>	!		<u>1MEQ/ML</u>	<u>A077394 002</u>	Nov 09, 2005
<u>AP</u>		HOSPIRA INC	<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>		HOSPIRA INC	1MEQ/ML	A202495 001	Mar 06, 2017

SODIUM CHLORIDE

INJECTABLE; INJECTION

BACTERIOSTATIC SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

<u>AP</u>		FRESENIUS KABI USA	<u>9MG/ML</u>	<u>A088911 001</u>	Feb 07, 1985
<u>AP</u>	+	!	<u>9MG/ML</u>	<u>N018800 001</u>	Oct 29, 1982

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>	+	!	<u>450MG/100ML</u>	<u>N018090 001</u>	

SODIUM CHLORIDE 0.9%

<u>AP</u>		SPECTRA MDCL DEVICES	<u>9MG/ML</u>	<u>A206171 001</u>	Jul 21, 2017
<u>AP</u>		WEST-WARD PHARMS INT	<u>9MG/ML</u>	<u>A201850 001</u>	Jan 20, 2012

SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

<u>AP</u>	+	!	<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>	+	BAXTER HLTHCARE	<u>9MG/ML</u>	<u>N016677 004</u>	Oct 30, 1985
<u>AP</u>	+		<u>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>9MG/ML</u>	<u>A088912 001</u>	Jan 10, 1985
<u>AP</u>			<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>	+	!	<u>9MG/ML</u>	<u>N018803 001</u>	Oct 29, 1982
<u>AP</u>	+		<u>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>	+		<u>900MG/100ML</u>	<u>N019480 001</u>	Sep 17, 1985
<u>AP</u>	+	!	<u>900MG/100ML</u>	<u>N016366 001</u>	
<u>AP</u>		ICU MEDICAL INC	<u>900MG/100ML</u>	<u>N016366 001</u>	
<u>AP</u>		JUBILANT CADISTA	<u>9MG/ML</u>	<u>A203352 001</u>	May 18, 2016
<u>AP</u>		LABORATORIOS GRIFOLS	<u>900MG/100ML</u>	<u>A207956 001</u>	May 25, 2017
<u>AP</u>	!	TARO	<u>9MG/ML</u>	<u>A077407 001</u>	Aug 11, 2006

PRESCRIPTION DRUG PRODUCT LIST

SODIUM CHLORIDE

INJECTABLE; INJECTION

SODIUM CHLORIDE 3% IN PLASTIC CONTAINER

<u>AP</u>	B BRAUN	<u>3GM/100ML</u>	<u>N019635</u>	<u>003</u>	Mar 09, 1988
<u>AP</u>	+! BAXTER HLTHCARE	<u>3GM/100ML</u>	<u>N019022</u>	<u>001</u>	Nov 01, 1983
<u>AP</u>	FRESENIUS KABI USA	<u>3GM/100ML</u>	<u>A209476</u>	<u>001</u>	Mar 13, 2019

SODIUM CHLORIDE 5% IN PLASTIC CONTAINER

<u>AP</u>	B BRAUN	<u>5GM/100ML</u>	<u>N019635</u>	<u>004</u>	Mar 09, 1988
<u>AP</u>	+ BAXTER HLTHCARE	<u>5GM/100ML</u>	<u>N019022</u>	<u>002</u>	Nov 01, 1983

SODIUM CHLORIDE 0.9%

+	B BRAUN	9MG/10ML	N019635	005	Aug 11, 2016
+	MEDEFIL INC	90MG/10ML (9MG/ML)	N202832	006	Jan 06, 2012
	WEST-WARD PHARMS	9MG/ML	A201833	001	Sep 24, 2013

INT

SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

+	LIEBEL-FLARSHEIM	1012.5MG/125ML (9MG/ML)	N021569	002	Jul 27, 2006
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SODIUM CHLORIDE IN PLASTIC CONTAINER

+	HOSPIRA	2.5MEQ/ML	N018897	001	Jul 20, 1984
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SOLUTION; IRRIGATION

SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

<u>AT</u>	+! B BRAUN	<u>900MG/100ML</u>	<u>N016733</u>	<u>001</u>	
<u>AT</u>	BAXTER HLTHCARE	<u>900MG/100ML</u>	<u>N017427</u>	<u>001</u>	
<u>AT</u>		<u>900MG/100ML</u>	<u>N017867</u>	<u>001</u>	
<u>AT</u>	ICU MEDICAL INC	<u>900MG/100ML</u>	<u>N017514</u>	<u>001</u>	
<u>AT</u>		<u>900MG/100ML</u>	<u>N018314</u>	<u>001</u>	

SOLUTION FOR SLUSH; IRRIGATION

SODIUM CHLORIDE 0.9% IN STERILE PLASTIC CONTAINER

	BAXTER HLTHCARE	900MG/100ML	N019319	002	May 17, 1985
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SODIUM FERRIC GLUCONATE COMPLEX

INJECTABLE; INJECTION

FERLECIIT

<u>AB</u>	+! SANOFI AVENTIS US	<u>62.5MG/5ML</u>	<u>N020955</u>	<u>001</u>	Feb 18, 1999
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SODIUM FERRIC GLUCONATE COMPLEX IN SUCROSE

<u>AB</u>	WEST-WARD PHARMS	<u>62.5MG/5ML</u>	<u>A078215</u>	<u>001</u>	Mar 31, 2011
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INT

SODIUM FLUORIDE F-18

INJECTABLE; INTRAVENOUS

SODIUM FLUORIDE F-18

<u>AP</u>	3D IMAGING DRUG	<u>10-200mCi/ML</u>	<u>A203777</u>	<u>001</u>	Oct 19, 2015
<u>AP</u>	BIOMEDCL RES FDN	<u>10-200mCi/ML</u>	<u>A204351</u>	<u>001</u>	Jan 09, 2015
<u>AP</u>	CARDINAL HEALTH 414	<u>10-200mCi/ML</u>	<u>A203780</u>	<u>001</u>	Jul 30, 2015
<u>AP</u>	DECATUR	<u>10-200mCi/ML</u>	<u>A204464</u>	<u>001</u>	Oct 21, 2014
<u>AP</u>	ESSENTIAL ISOTOPES	<u>10-200mCi/ML</u>	<u>A204541</u>	<u>001</u>	Oct 29, 2014
<u>AP</u>	HOT SHOTS NM LLC	<u>10-200mCi/ML</u>	<u>A204530</u>	<u>001</u>	Jul 29, 2015
<u>AP</u>	JUBILANT DRAXIMAGE	<u>10-200mCi/ML</u>	<u>A203968</u>	<u>001</u>	Oct 23, 2015
<u>AP</u>	KREITCHMAN PET CTR	<u>10-200mCi/ML</u>	<u>A203936</u>	<u>001</u>	May 19, 2016
<u>AP</u>	MIDWEST MEDCL	<u>10-200mCi/ML</u>	<u>A204440</u>	<u>001</u>	Nov 17, 2015
<u>AP</u>	MIPS CRF	<u>10-200mCi/ML</u>	<u>A204517</u>	<u>001</u>	Jul 21, 2015
<u>AP</u>	NCM USA BRONX LLC	<u>10-200mCi/ML</u>	<u>A204513</u>	<u>001</u>	Nov 28, 2014
<u>AP</u>	PETNET	<u>10-200mCi/ML</u>	<u>A203890</u>	<u>001</u>	Sep 28, 2015
<u>AP</u>	PRECISION NUCLEAR	<u>10-200mCi/ML</u>	<u>A204542</u>	<u>001</u>	Feb 27, 2015
<u>AP</u>	SHERTECH LABS LLC	<u>10-200mCi/ML</u>	<u>A204315</u>	<u>001</u>	Sep 22, 2014
<u>AP</u>	SOFIE	<u>10-200mCi/ML</u>	<u>A203592</u>	<u>001</u>	Aug 18, 2015
<u>AP</u>	!	<u>10-200mCi/ML</u>	<u>A203544</u>	<u>001</u>	Dec 26, 2012
<u>AP</u>	SPECTRON MRC LLC	<u>10-200mCi/ML</u>	<u>A203912</u>	<u>001</u>	Apr 22, 2015
<u>AP</u>	UCSF RODIOPHARM	<u>10-200mCi/ML</u>	<u>A204437</u>	<u>001</u>	Mar 13, 2014
<u>AP</u>	UNIV UTAH CYCLOTRON	<u>10-200mCi/ML</u>	<u>A204497</u>	<u>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

SODIUM IODIDE I-123

CAPSULE; ORAL

SODIUM IODIDE I 123

<u>AA</u>	+! CARDINAL HEALTH 418	<u>100uCi</u>	<u>N018671</u>	<u>001</u>	May 27, 1982
<u>AA</u>	+!	<u>200uCi</u>	<u>N018671</u>	<u>002</u>	May 27, 1982
<u>AA</u>	CURIUM	<u>100uCi</u>	<u>A071909</u>	<u>001</u>	Feb 28, 1989
<u>AA</u>		<u>200uCi</u>	<u>A071910</u>	<u>001</u>	Feb 28, 1989

PRESCRIPTION DRUG PRODUCT LIST

SODIUM IODIDE I-131

CAPSULE; ORAL

SODIUM IODIDE I 131

+ JUBILANT DRAXIMAGE 0.009-0.1mCi N021305 006 May 19, 2005

SOLUTION; ORAL

HICON**AA** +! JUBILANT DRAXIMAGE **250-1000mCi** **N021305 007** Dec 05, 2011**SODIUM IODIDE I 131****AA** INTL ISOTOPEs **250-1000mCi** **A209166 001** Feb 05, 2020SODIUM LACTATE

INJECTABLE; INJECTION

SODIUM LACTATE IN PLASTIC CONTAINER

+! HOSPIRA 5MEQ/ML N018947 001 Sep 05, 1984

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

NITHIODOTE

+! HOPE PHARMS 300MG/10ML (30MG/ML), N/A; N/A, 12.5GM/50ML (250MG/ML) N201444 001 Jan 14, 2011

SODIUM NITROPRUSSIDE

INJECTABLE; INJECTION

NITROPRESS**AP** ! HOSPIRA **25MG/ML** **A071961 001** Aug 01, 1988**SODIUM NITROPRUSSIDE****AP** AKORN **25MG/ML** **A208635 001** May 04, 2017**AP** AMNEAL **25MG/ML** **A209493 001** Nov 07, 2017**AP** AMPHASTAR PHARMS INC **25MG/ML** **A209832 001** Dec 18, 2017**AP** CAPLIN **25MG/ML** **A211016 001** Nov 29, 2019**AP** CIPLA **25MG/ML** **A210855 001** Jul 16, 2018**AP** DR REDDYS LABS LTD **25MG/ML** **A210114 001** Apr 10, 2019**AP** MEDICURE **25MG/ML** **A209584 001** Aug 10, 2018**AP** MICRO LABS **25MG/ML** **A209352 001** Dec 08, 2017**AP** MYLAN LABS LTD **25MG/ML** **A210763 001** Apr 17, 2018**AP** NEXUS PHARMS **25MG/ML** **A207499 001** May 25, 2017**AP** SAGENT PHARMS INC **25MG/ML** **A207426 001** Dec 08, 2016**AP** SOMERSET THERAPS LLC **25MG/ML** **A210882 001** Aug 17, 2018

SOLUTION; INTRAVENOUS

NIPRIDE RTU IN SODIUM CHLORIDE 0.9%

+! EXELA PHARMA SCS 20MG/100ML (0.2MG/ML) N209387 003 Jul 13, 2018

LLC

+! 50MG/100ML (0.5MG/ML) N209387 001 Mar 08, 2017

SODIUM OXYBATE

SOLUTION; ORAL

SODIUM OXYBATE**AA** HIKMA **500MG/ML** **A202090 001** Jan 17, 2017**XYREM****AA** +! JAZZ PHARMS **500MG/ML** **N021196 001** Jul 17, 2002SODIUM PHENYLBUTYRATE

POWDER; ORAL

BUPHENYL**AB** +! HORIZON THERAP **3GM/TEASPOONFUL** **N020573 001** Apr 30, 1996**SODIUM PHENYLBUTYRATE****AB** PAR PHARM **3GM/TEASPOONFUL** **A203918 001** Jun 15, 2016**AB** SIGMAPHARM LABS LLC **3GM/TEASPOONFUL** **A202819 001** Mar 22, 2013

TABLET; ORAL

BUPHENYL**AB** +! HORIZON THERAP **500MG** **N020572 001** May 13, 1996**SODIUM PHENYLBUTYRATE****AB** PAR PHARM **500MG** **A204395 001** Apr 15, 2016

PRESCRIPTION DRUG PRODUCT LIST

SODIUM PHOSPHATE, DIBASIC, ANHYDROUS; SODIUM PHOSPHATE, MONOBASIC, MONOHYDRATE

TABLET; ORAL

OSMOPREP

+! SALIX PHARMS 0.398GM;1.102GM N021892 001 Mar 16, 2006

SODIUM PHOSPHATE, DIBASIC, HEPTAHYDRATE; SODIUM PHOSPHATE, MONOBASIC, ANHYDROUS

INJECTABLE; INJECTION

SODIUM PHOSPHATES IN PLASTIC CONTAINER

+! HOSPIRA 142MG/ML;276MG/ML N018892 001 May 10, 1983

SODIUM POLYSTYRENE SULFONATE

POWDER; ORAL, RECTAL

KALEXATE**AA** ! KVK TECH **454GM/BOT** **A040905 001** Mar 30, 2009**KIONEX****AA** PADDOCK LLC **454GM/BOT** **A040029 001** Feb 06, 1998**SODIUM POLYSTYRENE SULFONATE****AA** BELCHER **454GM/BOT** **A205727 001** Feb 23, 2016**AA** CMP PHARMA INC **454GM/BOT** **A089910 001** Jan 19, 1989**AA** ECI PHARMS LLC **453.6GM/BOT** **A090313 001** Dec 21, 2011**AA** EPIC PHARMA LLC **453.6GM/BOT** **A202333 001** Mar 19, 2014**AA** NOVELGENIX THERAPS **454GM/BOT** **A206815 001** Feb 18, 2016**AA** NUVO PHARMS INC **454GM/BOT** **A204071 001** Nov 28, 2014

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** PADDOCK LLC **15GM/60ML** **A040028 001** Sep 17, 2007**SODIUM POLYSTYRENE SULFONATE****AA** PADDOCK LLC **15GM/60ML** **A090590 001** May 13, 2011**SPS****AA** ! CMP PHARMA INC **15GM/60ML** **A087859 001** Dec 08, 1982SODIUM TETRADECYL SULFATE

INJECTABLE; INJECTION

SODIUM TETRADECYL SULFATE**AP** CUSTOPHARM INC **60MG/2ML (30MG/ML)** **A209937 001** Dec 09, 2019**SOTRADECOL****AP** ! MYLAN INSTITUTIONAL **60MG/2ML (30MG/ML)** **A040541 002** Nov 12, 2004

! 20MG/2ML (10MG/ML) A040541 001 Nov 12, 2004

SODIUM THIOSULFATE

SOLUTION; INTRAVENOUS

SODIUM THIOSULFATE

+! HOPE PHARMS 12.5GM/50ML (250MG/ML) N203923 001 Feb 14, 2012

SODIUM ZIRCONIUM CYCLOSILICATE

FOR SUSPENSION; ORAL

LOKELMA

+ ASTRAZENECA PHARMS

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

TABLET; ORAL

EPCLUSA

+! GILEAD SCIENCES INC 400MG;100MG

N208341 001 Jun 28, 2016

SOFOBUVIR; VELPATASVIR; VOXILAPREVIR

TABLET; ORAL

VOSEVI

+! GILEAD SCIENCES INC 400MG;100MG;100MG

N209195 001 Jul 18, 2017

PRESCRIPTION DRUG PRODUCT LISTSOLIFENACIN SUCCINATE

TABLET; ORAL

SOLIFENACIN SUCCINATE

AB	AJANTA PHARMA LTD	5MG	A205483 001	May 20, 2019
AB		10MG	A205483 002	May 20, 2019
AB	ALEMBIC PHARMS LTD	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	AMNEAL PHARMS CO	5MG	A209719 001	May 20, 2019
AB		10MG	A209719 002	May 20, 2019
AB	BRECKENRIDGE	5MG	A209818 001	May 20, 2019
AB		10MG	A209818 002	May 20, 2019
AB	CELLTRION	5MG	A210582 001	May 20, 2019
AB		10MG	A210582 002	May 20, 2019
AB	CIPLA	5MG	A209839 001	May 20, 2019
AB		10MG	A209839 002	May 20, 2019
AB	CSPC OUYI	5MG	A211423 001	Dec 11, 2019
AB		10MG	A211423 002	Dec 11, 2019
AB	GLENMARK PHARMS INC	5MG	A209239 001	May 20, 2019
AB		10MG	A209239 002	May 20, 2019
AB	JIANGXI BOYA SEEHOT	5MG	A210281 001	May 20, 2019
AB		10MG	A210281 002	May 20, 2019
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	SCIEGEN PHARMS INC	5MG	A211657 001	May 20, 2019
AB		10MG	A211657 002	May 20, 2019
AB	STRIDES PHARMA	5MG	A212214 001	Sep 26, 2019
AB		10MG	A212214 002	Sep 26, 2019
AB	TEVA PHARMS USA	5MG	A091464 001	Apr 02, 2014
AB		10MG	A091464 002	Apr 02, 2014
AB	UNICHEM LABS LTD	5MG	A211701 001	Aug 27, 2019
AB		10MG	A211701 002	Aug 27, 2019
AB	WATSON LABS INC	5MG	A202551 001	May 20, 2019
AB		10MG	A202551 002	May 20, 2019
<u>VESICARE</u>				
AB	+ ASTELLAS	5MG	N021518 001	Nov 19, 2004
AB	+!	10MG	N021518 002	Nov 19, 2004

SOLRIAMFETOL HYDROCHLORIDE

TABLET; ORAL

SUNOSI

+	JAZZ	EQ 75MG BASE	N211230 001	Jun 17, 2019
+		EQ 150MG BASE	N211230 002	Jun 17, 2019

SOMATROPIN

INJECTABLE; INJECTION

GENOTROPIN

BX	+! PHARMACIA	5.8MG/VIAL	N020280 006	Aug 24, 1995
HUMATROPE				
BX	+! LILLY	5MG/VIAL	N019640 004	Mar 08, 1987
BX	+	6MG/VIAL	N019640 005	Feb 04, 1999
NORDITROPIN FLEXPRO				
BX	NOVO NORDISK INC	5MG/1.5ML	N021148 008	Mar 01, 2010
BX		10MG/1.5ML	N021148 009	Mar 01, 2010
OMNITROPE				
BX	SANDOZ	1.5MG/VIAL	N021426 002	May 30, 2006
BX		5MG/1.5ML	N021426 003	Jan 16, 2008
BX		5.8MG/VIAL	N021426 001	May 30, 2006
BX		10MG/1.5ML	N021426 004	Aug 25, 2008
SAIZEN				
BX	+ EMD SERONO	5MG/VIAL	N019764 002	Oct 08, 1996
SEROSTIM				
BX	EMD SERONO	5MG/VIAL	N020604 002	Aug 23, 1996
BX		6MG/VIAL	N020604 001	Aug 23, 1996
ZOMACTON				
BX	+! FERRING	5MG/VIAL	N019774 002	Jan 04, 2002
GENOTROPIN				
	+! PHARMACIA	13.8MG/VIAL	N020280 007	Oct 23, 1996
GENOTROPIN PRESERVATIVE FREE				
	+ PHARMACIA	0.2MG/VIAL	N020280 001	Jan 27, 1998

PRESCRIPTION DRUG PRODUCT LISTSOMATROPIN

INJECTABLE; INJECTION

GENOTROPIN PRESERVATIVE FREE

+		0.4MG/VIAL	N020280 002	Jan 27, 1998
+		0.6MG/VIAL	N020280 003	Jan 27, 1998
+		0.8MG/VIAL	N020280 005	Jan 27, 1998
+		1MG/VIAL	N020280 008	Jan 27, 1998
+		1.2MG/VIAL	N020280 009	Jan 27, 1998
+		1.4MG/VIAL	N020280 010	Jan 27, 1998
+		1.6MG/VIAL	N020280 011	Jan 27, 1998
+		1.8MG/VIAL	N020280 012	Jan 27, 1998
+	!	2MG/VIAL	N020280 013	Jan 27, 1998

HUMATROPE

+	!	LILLY	12MG/VIAL	N019640 006	Feb 04, 1999
+	!		24MG/VIAL	N019640 007	Feb 04, 1999

NORDITROPIN FLEXPRO

		NOVO NORDISK INC	15MG/1.5ML	N021148 010	Mar 01, 2010
			30MG/3ML	N021148 011	Jan 23, 2015

NUTROPIN AQ NUSPIN

+	!	GENENTECH	5MG/2ML (2.5MG/ML)	N020522 003	Jan 03, 2008
+	!		10MG/2ML (5MG/ML)	N020522 005	Jan 03, 2008
+	!		20MG/2ML (10MG/ML)	N020522 004	Jan 03, 2008

SAIZEN

+	!	EMD SERONO	8.8MG/VIAL	N019764 003	Aug 29, 2000
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SEROSTIM

		EMD SERONO	4MG/VIAL	N020604 003	Jul 25, 1997
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ZOMACTON

+		FERRING	10MG/VIAL	N019774 003	Mar 07, 2012
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SOMATROPIN RECOMBINANT

INJECTABLE; INJECTION

ZORBTIVE

+	!	EMD SERONO	8.8MG/VIAL	N021597 004	Dec 01, 2003
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SONIDEGIB PHOSPHATE

CAPSULE; ORAL

ODOMZO

+	!	SUN PHARMA GLOBAL	EQ 200MG BASE	N205266 001	Jul 24, 2015
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SORAFENIB TOSYLATE

TABLET; ORAL

NEXAVAR

+	!	BAYER HLTHCARE	EQ 200MG BASE	N021923 001	Dec 20, 2005
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SORBITOL

SOLUTION; IRRIGATION

SORBITOL 3% IN PLASTIC CONTAINER

		BAXTER HLTHCARE	3GM/100ML	N017863 001	
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SORBITOL 3.3% IN PLASTIC CONTAINER

		B BRAUN	3.3GM/100ML	N016741 001	
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SOTALOL HYDROCHLORIDE

SOLUTION; INTRAVENOUS

SOTALOL HYDROCHLORIDE

+	!	ALTATHERA PHARMS LLC	150MG/10ML (15MG/ML)	N022306 001	Jul 02, 2009
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SOLUTION; ORAL

SOTYLIZE

+	!	ARBOR PHARMS LLC	5MG/ML (5MG/ML)	N205108 001	Oct 22, 2014
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TABLET; ORAL

BETAPACE

<u>AB</u>	+	COVIS PHARMA BV	<u>80MG</u>	<u>N019865 001</u>	Oct 30, 1992
<u>AB</u>	+		<u>120MG</u>	<u>N019865 005</u>	Apr 20, 1994
<u>AB</u>	+	!	<u>160MG</u>	<u>N019865 002</u>	Oct 30, 1992
<u>AB</u>	+		<u>240MG</u>	<u>N019865 003</u>	Oct 30, 1992

BETAPACE AF

<u>AB</u>	+	COVIS PHARMA BV	<u>80MG</u>	<u>N021151 001</u>	Feb 22, 2000
<u>AB</u>	+		<u>120MG</u>	<u>N021151 002</u>	Feb 22, 2000
<u>AB</u>	+	!	<u>160MG</u>	<u>N021151 003</u>	Feb 22, 2000

SORINE

<u>AB</u>		UPSHER SMITH LABS	<u>80MG</u>	<u>A075500 001</u>	Apr 27, 2001
<u>AB</u>			<u>120MG</u>	<u>A075500 004</u>	Apr 27, 2001
<u>AB</u>			<u>160MG</u>	<u>A075500 002</u>	Apr 27, 2001
<u>AB</u>			<u>240MG</u>	<u>A075500 003</u>	Apr 27, 2001

PRESCRIPTION DRUG PRODUCT LIST

SPIRONOLACTONE

TABLET; ORAL

SPIRONOLACTONE

AB		100MG	A040750 003	Aug 29, 2006
AB	SUN PHARM INDUSTRIES	25MG	A089424 001	Jul 23, 1986
AB		50MG	A089424 002	Aug 11, 1999
AB		100MG	A089424 003	Aug 11, 1999
AB	ZYDUS PHARMS	25MG	A205936 001	Jul 18, 2018
AB		50MG	A205936 002	Jul 18, 2018
AB		100MG	A205936 003	Jul 18, 2018

STAVUDINE

CAPSULE; ORAL

STAVUDINE

AB	AUROBINDO PHARMA	15MG	A077672 003	Dec 29, 2008
AB		20MG	A077672 004	Dec 29, 2008
AB		30MG	A077672 001	Dec 29, 2008
AB		40MG	A077672 002	Dec 29, 2008

ZERIT

AB	+ BRISTOL MYERS SQUIBB	15MG	N020412 002	Jun 24, 1994
AB	+	20MG	N020412 003	Jun 24, 1994
AB	+	30MG	N020412 004	Jun 24, 1994
AB	+!	40MG	N020412 005	Jun 24, 1994

STERILE WATER FOR INJECTION

LIQUID; N/A

BACTERIOSTATIC WATER FOR INJECTION IN PLASTIC CONTAINER

AP	+! HOSPIRA	100%	N018802 001	Oct 27, 1982
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STERILE WATER FOR INJECTION

AP	FRESENIUS KABI USA	100%	A209689 001	Nov 24, 2017
AP	+! HOSPIRA	100% (10ML)	N018801 002	Oct 27, 1982
AP	MEDEFIL INC	100% (10ML)	A211188 005	Dec 02, 2019
AP	WEST-WARD PHARMS INT	100% (10ML)	A206369 001	Sep 02, 2015

STERILE WATER FOR INJECTION IN PLASTIC CONTAINER

AP	+! B BRAUN	100%	N019633 001	Feb 29, 1988
AP	+! BAXTER HLTHCARE	100%	N018632 001	Jun 30, 1982
AP	+!	100%	N018632 002	Apr 19, 1988
AP	FRESENIUS KABI USA	100%	A088400 001	Jan 16, 1984
AP	+! ICU MEDICAL INC	100%	N018233 001	
AP	+!	100%	N019869 001	Dec 26, 1989
AP	TARO	100%	A077393 001	Aug 11, 2006

STERILE WATER FOR INJECTION

+!	HOSPIRA	100% (20ML)	N018801 003	Oct 27, 1982
+!		100% (50ML)	N018801 004	Oct 27, 1982
+!		100% (100ML)	N018801 005	Oct 27, 1982
	MEDEFIL INC	100% (1ML)	A211188 001	Dec 02, 2019
!		100% (2.5ML)	A211188 002	Dec 02, 2019
!		100% (3ML)	A211188 003	Dec 02, 2019
!		100% (5ML)	A211188 004	Dec 02, 2019

STERILE WATER FOR IRRIGATION

LIQUID; IRRIGATION

STERILE WATER

AT	+ BAXTER HLTHCARE	100%	N017428 001	
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STERILE WATER IN PLASTIC CONTAINER

AT	+ B BRAUN	100%	N016734 001	
AT	BAXTER HLTHCARE	100%	N017866 001	
AT	ICU MEDICAL INC	100%	N017513 001	
AT		100%	N018313 001	

STIRIPENTOL

CAPSULE; ORAL

DIACOMIT

+	BIOCODEX SA	250MG	N206709 001	Aug 20, 2018
+	!	500MG	N206709 002	Aug 20, 2018

FOR SUSPENSION; ORAL

DIACOMIT

+	BIOCODEX SA	250MG/PACKET	N207223 001	Aug 20, 2018
+	!	500MG/PACKET	N207223 002	Aug 20, 2018

PRESCRIPTION DRUG PRODUCT LIST

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 INC**20MG/ML****N008453 002**QUELICIN**AP** +! HOSPIRA**20MG/ML****N008845 006**SUCCINYLCHOLINE CHLORIDE**AP** AMNEAL**20MG/ML****A211432 001** Nov 16, 2018**AP** AMRING PHARMS**20MG/ML****A210231 001** Jun 04, 2018**AP** BRECKENRIDGE**20MG/ML****A212638 001** Oct 09, 2019**AP** DR REDDYS**20MG/ML****A210698 001** Aug 02, 2019**AP** SOMERSET THERAPS**20MG/ML****A211589 001** Jan 15, 2020

LLC

AP ! ZYDUS PHARMS**20MG/ML****A209467 001** May 04, 2018SUCRALFATE

SUSPENSION; ORAL

CARAFATE**AB** +! ALLERGAN**1GM/10ML****N019183 001** Dec 16, 1993SUCRALFATE**AB** AMNEAL PHARMS LLC**1GM/10ML****A209356 001** Dec 02, 2019

TABLET; ORAL

CARAFATE**AB** +! ALLERGAN**1GM****N018333 001**SUCRALFATE**AB** TEVA**1GM****A070848 001** Mar 29, 1996SUCROFERRIC OXYHYDROXIDE

TABLET, CHEWABLE; ORAL

VELPHORO

+! VIFOR FRESENIUS

500MG

N205109 001 Nov 27, 2013

SUFENTANIL CITRATE

INJECTABLE; INJECTION

SUFENTA PRESERVATIVE FREE**AP** +! AKORN**EQ 0.05MG BASE/ML****N019050 001** May 04, 1984SUFENTANIL CITRATE**AP** HOSPIRA**EQ 0.05MG BASE/ML****A074534 001** Dec 11, 1996**AP** WEST-WARD PHARMS**EQ 0.05MG BASE/ML****A074413 001** Dec 15, 1995

INT

TABLET; SUBLINGUAL

DSUVIA

+! ACELRX PHARMS

EQ 0.03MG BASE

N209128 001 Nov 02, 2018

SUGAMMADEX SODIUM

SOLUTION; INTRAVENOUS

BRIDION

+ ORGANON SUB MERCK

EQ 200MG BASE/2ML (EQ 100MG BASE/ML)

N022225 002 Dec 15, 2015

+!

EQ 500MG BASE/5ML (EQ 100MG BASE/ML)

N022225 001 Dec 15, 2015

PRESCRIPTION DRUG PRODUCT LIST

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 +!** VALEANT PHARMS
NORTH**10%****N019931 001** Dec 23, 1996**SULFACETAMIDE SODIUM****AB** FOUGERA PHARMS**10%****A077015 001** Nov 17, 2006**AB** PERRIGO CO**10%****A078649 001** Mar 23, 2009

TENNESSEE

AB TARO**10%****A078668 001** May 20, 2009

OINTMENT; OPHTHALMIC

SULFACETAMIDE SODIUM

! PERRIGO CO

10%

A080029 001

TENNESSEE

SOLUTION/DROPS; OPHTHALMIC

BLEPH-10**AT !** ALLERGAN**10%****A080028 001****SULFACETAMIDE SODIUM****AT** AKORN**10%****A040215 001** May 25, 1999**AT** BAUSCH AND LOMB**10%****A040066 001** Dec 28, 1994**AT** SANDOZ INC**10%****A089560 001** Oct 18, 1988SULFADIAZINE

TABLET; ORAL

SULFADIAZINE

! SANDOZ

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

SUSPENSION; ORAL

SULFAMETHOXAZOLE AND TRIMETHOPRIM**AB** ANI PHARMS INC**200MG/5ML; 40MG/5ML****A077612 001** Nov 13, 2006**AB** AUROBINDO PHARMA**200MG/5ML; 40MG/5ML****A091348 001** Jun 08, 2010**AB !** HI TECH PHARMA**200MG/5ML; 40MG/5ML****A074650 001** Dec 29, 1997**AB** LANNETT CO INC**200MG/5ML; 40MG/5ML****A077785 001** Jan 24, 2007**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****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** AUROBINDO PHARMA**800MG; 160MG****A076899 002** Jan 27, 2005**AB** AUROBINDO PHARMA**400MG; 80MG****A090624 001** Feb 16, 2010**AB** AUROBINDO PHARMA**800MG; 160MG****A090624 002** Feb 16, 2010**AB** CHARTWELL MOLECULES**400MG; 80MG****A078060 002** Jan 25, 2007**AB** CHARTWELL MOLECULES**800MG; 160MG****A078060 001** Jan 25, 2007**AB** GLENMARK GENERICS**400MG; 80MG****A090828 002** Dec 22, 2010**AB** GLENMARK GENERICS**800MG; 160MG****A090828 001** Dec 22, 2010**AB** SUN PHARM**400MG; 80MG****A071017 002** Aug 25, 1986

INDUSTRIES

AB SUN PHARM**800MG; 160MG****A071017 001** Aug 25, 1986**AB** VISTA PHARMS**400MG; 80MG****A076817 001** Oct 07, 2005**AB** VISTA PHARMS**800MG; 160MG****A076817 002** Oct 07, 2005

PRESCRIPTION DRUG PRODUCT LIST

SULFANILAMIDECREAM; VAGINAL
AVC

+! MYLAN SPECIALITY LP 15% N006530 003 Jan 27, 1987

SULFASALAZINE

TABLET; ORAL

AZULFIDINE**AB** +! PHARMACIA AND UPJOHN **500MG** **N007073 001**SULFASALAZINE**AB** VINTAGE PHARMS **500MG** **A040349 001** Jan 11, 2002
AB WATSON LABS **500MG** **A085828 001**

TABLET, DELAYED RELEASE; ORAL

AZULFIDINE EN-TABS**AB** +! PHARMACIA AND UPJOHN **500MG** **N007073 002** Apr 06, 1983SULFASALAZINE**AB** VINTAGE PHARMS **500MG** **A075339 001** Jan 11, 2002SULFUR HEXAFLUORIDE LIPID-TYPE A MICROSPHERESFOR 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** MYLAN **150MG** **A073039 002** Jun 22, 1993**AB** **200MG** **A073039 001** Jun 22, 1993**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, 1990SUMATRIPTAN

SPRAY; NASAL

IMITREX**AB** +! GLAXOSMITHKLINE **5MG/SPRAY** **N020626 001** Aug 26, 1997**AB** +! **20MG/SPRAY** **N020626 003** Aug 26, 1997SUMATRIPTAN**AB** LANNETT CO INC **5MG/SPRAY** **A204841 001** Feb 19, 2016**AB** **20MG/SPRAY** **A204841 002** Feb 19, 2016

TOSYMRA

+! UPSHER SMITH LABS 10MG/SPRAY N210884 001 Jan 25, 2019

SUMATRIPTAN SUCCINATE

INJECTABLE; SUBCUTANEOUS

IMITREX STATDOSE**AB** +! GLAXOSMITHKLINE **EQ 4MG BASE/0.5ML (EQ 8MG BASE/ML)** **N020080 002** Feb 01, 2006**AB** +! **EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)** **N020080 003** Dec 23, 1996SUMATRIPTAN SUCCINATE**AB** ANTARES PHARMA INC **EQ 4MG BASE/0.5ML (EQ 8MG BASE/ML)** **A078319 001** Dec 10, 2015**AB** **EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)** **A078319 002** Dec 10, 2015**AB** DR REDDYS **EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)** **A090495 001** Jan 29, 2014**AB** SUN PHARM **EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)** **A090358 001** Jun 21, 2011IMITREX**AP** +! GLAXOSMITHKLINE **EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)** **N020080 001** Dec 28, 1992SUMATRIPTAN SUCCINATE**AP** AUROBINDO PHARMA LTD **EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)** **A202758 001** Apr 23, 2013**AP** FRESenius KABI USA **EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)** **A079242 001** Mar 02, 2009**AP** HIKMA **EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)** **A200183 001** Sep 16, 2013**AP** MYLAN ASI **EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)** **A090314 001** Jun 10, 2010**AP** **EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)** **A090641 001** Jul 28, 2010**AP** PAR PHARM **EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)** **A077332 001** Oct 09, 2009**AP** WEST-WARD PHARMS INT **EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)** **A079123 001** Feb 06, 2009**AP** WOCKHARDT **EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)** **A078593 001** Feb 06, 2009

POWDER; NASAL

ONZETRA XSAIL

+! CURRAX EQ 11MG BASE N206099 001 Jan 27, 2016

PRESCRIPTION DRUG PRODUCT LIST

SUMATRIPTAN SUCCINATE

SOLUTION;SUBCUTANEOUS

ZEMBRACE SYMTOUCH

+ UPSHER SMITH LABS

EQ 3MG BASE/0.5ML (EQ 3MG BASE/0.5ML)

N208223 001 Jan 28, 2016

TABLET;ORAL

IMITREXAB + GLAXOSMITHKLINEEQ 25MG BASEN020132 002 Jun 01, 1995AB +EQ 50MG BASEN020132 003 Jun 01, 1995AB +!EQ 100MG BASEN020132 001 Jun 01, 1995SUMATRIPTAN SUCCINATEAB AUROBINDO PHARMAEQ 25MG BASEA078327 001 Aug 10, 2009ABEQ 50MG BASEA078327 002 Aug 10, 2009ABEQ 100MG BASEA078327 003 Aug 10, 2009AB COREPHARMAEQ 25MG BASEA200263 001 Jun 19, 2012ABEQ 50MG BASEA200263 002 Jun 19, 2012ABEQ 100MG BASEA200263 003 Jun 19, 2012AB DR REDDYS LABS INCEQ 25MG BASEA076847 001 Aug 10, 2009ABEQ 50MG BASEA076847 002 Aug 10, 2009ABEQ 100MG BASEA076847 003 Aug 10, 2009AB MYLANEQ 25MG BASEA077744 001 Aug 10, 2009ABEQ 50MG BASEA077744 002 Aug 10, 2009ABEQ 100MG BASEA077744 003 Aug 10, 2009AB ORCHID HLTHCAREEQ 25MG BASEA078284 001 Aug 10, 2009ABEQ 50MG BASEA078284 002 Aug 10, 2009ABEQ 100MG BASEA078284 003 Aug 10, 2009AB SUN PHARM INDSEQ 25MG BASEA078295 001 Aug 10, 2009ABEQ 50MG BASEA078295 002 Aug 10, 2009ABEQ 100MG BASEA078295 003 Aug 10, 2009AB SUN PHARM INDS LTDEQ 25MG BASEA076554 001 Aug 10, 2009ABEQ 50MG BASEA076554 002 Aug 10, 2009ABEQ 100MG BASEA076572 001 Feb 09, 2009AB WATSON LABSEQ 25MG BASEA076933 001 Aug 10, 2009ABEQ 50MG BASEA076933 002 Aug 10, 2009ABEQ 100MG BASEA076933 003 Aug 10, 2009SUNITINIB MALATE

CAPSULE;ORAL

SUTENT

+ CPPI CV

EQ 12.5MG BASE

N021938 001 Jan 26, 2006

+

EQ 25MG BASE

N021938 002 Jan 26, 2006

+

EQ 37.5MG BASE

N021938 004 Mar 31, 2009

+!

EQ 50MG BASE

N021938 003 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

PROGRAFAB + ASTELLASEQ 0.5MG BASEN050708 003 Aug 24, 1998AB +EQ 1MG BASEN050708 001 Apr 08, 1994AB +!EQ 5MG BASEN050708 002 Apr 08, 1994TACROLIMUSAB ACCORD HLTHCAREEQ 0.5MG BASEA091195 001 Aug 31, 2011ABEQ 1MG BASEA091195 002 Aug 31, 2011ABEQ 5MG BASEA091195 003 Aug 31, 2011AB BELCHEREQ 0.5MG BASEA206651 001 Nov 30, 2017ABEQ 1MG BASEA206651 002 Nov 30, 2017ABEQ 5MG BASEA206651 003 Nov 30, 2017AB DR REDDYS LABS LTDEQ 0.5MG BASEA090509 001 May 12, 2010ABEQ 1MG BASEA090509 002 May 12, 2010ABEQ 5MG BASEA090509 003 May 12, 2010AB MYLANEQ 0.5MG BASEA090596 001 Sep 17, 2010ABEQ 1MG BASEA090596 002 Sep 17, 2010ABEQ 5MG BASEA090596 003 Sep 17, 2010AB PANACEA BIOTEC LTDEQ 0.5MG BASEA090802 001 Sep 28, 2012ABEQ 1MG BASEA090802 002 Sep 28, 2012ABEQ 5MG BASEA090802 003 Sep 28, 2012AB SANDOZEQ 0.5MG BASEA065461 001 Aug 10, 2009

PRESCRIPTION DRUG PRODUCT LIST

TACROLIMUS

CAPSULE; ORAL

TACROLIMUS

AB		<u>EQ 1MG BASE</u>	<u>A065461 002</u>	Aug 10, 2009
AB		<u>EQ 5MG BASE</u>	<u>A065461 003</u>	Aug 10, 2009
AB	STRIDES PHARMA	<u>EQ 0.5MG BASE</u>	<u>A090687 001</u>	Jul 22, 2014
AB		<u>EQ 1MG BASE</u>	<u>A090687 002</u>	Jul 22, 2014
AB		<u>EQ 5MG BASE</u>	<u>A090687 003</u>	Jul 22, 2014

CAPSULE, EXTENDED RELEASE; ORAL

ASTAGRAF XL

+	ASTELLAS	EQ 0.5MG BASE	N204096 001	Jul 19, 2013
+		EQ 1MG BASE	N204096 002	Jul 19, 2013
+	!	EQ 5MG BASE	N204096 003	Jul 19, 2013

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

AP	+	ASTELLAS	<u>EQ 5MG BASE/ML</u>	<u>N050709 001</u>	Apr 08, 1994
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TACROLIMUS

AP		HOSPIRA INC	<u>EQ 5MG BASE/ML</u>	<u>A203900 001</u>	Aug 25, 2017
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OINTMENT; TOPICAL

PROTOPIC

AB	+	LEO PHARMA AS	<u>0.03%</u>	<u>N050777 001</u>	Dec 08, 2000
AB	+	!	<u>0.1%</u>	<u>N050777 002</u>	Dec 08, 2000

TACROLIMUS

AB		ACCORD HLTHCARE	<u>0.03%</u>	<u>A211688 001</u>	Jan 31, 2019
AB			<u>0.1%</u>	<u>A211688 002</u>	Jan 31, 2019
AB		FOUGERA PHARMS INC	<u>0.03%</u>	<u>A200744 001</u>	Sep 09, 2014
AB			<u>0.1%</u>	<u>A200744 002</u>	Sep 09, 2014
AB		GLENMARK PHARMS LTD	<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

TABLET; ORAL

CIALIS

AB	+	LILLY	<u>2.5MG</u>	<u>N021368 004</u>	Jan 07, 2008
AB	+		<u>5MG</u>	<u>N021368 001</u>	Nov 21, 2003
AB	+		<u>10MG</u>	<u>N021368 002</u>	Nov 21, 2003

TADALAFIL

AB		ACCORD HLTHCARE	<u>2.5MG</u>	<u>A209167 001</u>	Mar 26, 2019
AB			<u>5MG</u>	<u>A209167 002</u>	Mar 26, 2019
AB		AJANTA PHARMA LTD	<u>2.5MG</u>	<u>A209654 001</u>	Mar 26, 2019
AB			<u>5MG</u>	<u>A209654 002</u>	Mar 26, 2019
AB			<u>10MG</u>	<u>A209654 003</u>	Mar 26, 2019
AB		ALEMBIC PHARMS LTD	<u>2.5MG</u>	<u>A204809 001</u>	Mar 26, 2019
AB			<u>5MG</u>	<u>A204809 002</u>	Mar 26, 2019
AB			<u>10MG</u>	<u>A204809 003</u>	Mar 26, 2019
AB		AMNEAL PHARMS CO	<u>2.5MG</u>	<u>A209744 001</u>	Mar 26, 2019
AB			<u>5MG</u>	<u>A209744 002</u>	Mar 26, 2019
AB			<u>10MG</u>	<u>A209744 003</u>	Mar 26, 2019
AB		AUROBINDO PHARMA LTD	<u>2.5MG</u>	<u>A206285 001</u>	Mar 26, 2019
AB			<u>5MG</u>	<u>A206285 002</u>	Mar 26, 2019
AB			<u>10MG</u>	<u>A206285 003</u>	Mar 26, 2019
AB		CIPLA	<u>2.5MG</u>	<u>A209539 001</u>	Mar 26, 2019
AB			<u>5MG</u>	<u>A209539 002</u>	Mar 26, 2019
AB			<u>10MG</u>	<u>A209539 003</u>	Mar 26, 2019
AB		DR REDDYS LABS LTD	<u>2.5MG</u>	<u>A210069 001</u>	Mar 26, 2019
AB			<u>5MG</u>	<u>A210069 002</u>	Mar 26, 2019
AB			<u>10MG</u>	<u>A210069 003</u>	Mar 26, 2019
AB		HETERO LABS LTD III	<u>2.5MG</u>	<u>A209908 001</u>	Mar 26, 2019
AB			<u>5MG</u>	<u>A209908 002</u>	Mar 26, 2019
AB			<u>10MG</u>	<u>A209908 003</u>	Mar 26, 2019
AB		LUPIN LTD	<u>2.5MG</u>	<u>A210567 001</u>	Mar 26, 2019
AB			<u>5MG</u>	<u>A210567 002</u>	Mar 26, 2019
AB			<u>10MG</u>	<u>A210567 003</u>	Mar 26, 2019
AB		MACLEODS PHARMS LTD	<u>2.5MG</u>	<u>A207244 001</u>	Oct 07, 2019

PRESCRIPTION DRUG PRODUCT LISTTADALAFIL

TABLET; ORAL

TADALAFIL

<u>AB</u>		<u>5MG</u>	<u>A207244 002</u>	Oct 07, 2019
<u>AB</u>		<u>10MG</u>	<u>A207244 003</u>	Oct 07, 2019
<u>AB</u>	MYLAN	<u>2.5MG</u>	<u>A206956 001</u>	Apr 29, 2019
<u>AB</u>		<u>5MG</u>	<u>A206957 001</u>	Apr 29, 2019
<u>AB</u>		<u>10MG</u>	<u>A206956 002</u>	Apr 29, 2019
<u>AB</u>	QILU	<u>2.5MG</u>	<u>A210420 001</u>	Mar 26, 2019
<u>AB</u>		<u>5MG</u>	<u>A210420 002</u>	Mar 26, 2019
<u>AB</u>		<u>10MG</u>	<u>A210420 003</u>	Mar 26, 2019
<u>AB</u>	SUN PHARM	<u>2.5MG</u>	<u>A208934 001</u>	Mar 26, 2019
<u>AB</u>		<u>5MG</u>	<u>A208934 002</u>	Mar 26, 2019
<u>AB</u>		<u>10MG</u>	<u>A208934 003</u>	Mar 26, 2019
<u>AB</u>	SUNSHINE LAKE	<u>2.5MG</u>	<u>A211335 001</u>	Mar 26, 2019
<u>AB</u>		<u>5MG</u>	<u>A211335 002</u>	Mar 26, 2019
<u>AB</u>		<u>10MG</u>	<u>A211335 003</u>	Mar 26, 2019
<u>AB</u>	TEVA PHARMS USA	<u>2.5MG</u>	<u>A090141 001</u>	May 22, 2018
<u>AB</u>		<u>5MG</u>	<u>A090141 002</u>	May 22, 2018
<u>AB</u>		<u>10MG</u>	<u>A090141 003</u>	May 22, 2018
<u>AB</u>	TORRENT	<u>2.5MG</u>	<u>A211839 001</u>	Mar 26, 2019
<u>AB</u>		<u>5MG</u>	<u>A211839 002</u>	Mar 26, 2019
<u>AB</u>		<u>10MG</u>	<u>A211839 003</u>	Mar 26, 2019
<u>AB</u>	UNICHEM LABS LTD	<u>2.5MG</u>	<u>A209250 001</u>	Mar 26, 2019
<u>AB</u>		<u>5MG</u>	<u>A209250 002</u>	Mar 26, 2019
<u>AB</u>		<u>10MG</u>	<u>A209250 003</u>	Mar 26, 2019
<u>AB</u>	ZYDUS PHARMS	<u>2.5MG</u>	<u>A206693 001</u>	Mar 26, 2019
<u>AB</u>		<u>5MG</u>	<u>A206693 002</u>	Mar 26, 2019
<u>AB</u>		<u>10MG</u>	<u>A206693 003</u>	Mar 26, 2019

CIALIS

<u>AB1</u>	+!	LILLY	<u>20MG</u>	<u>N021368 003</u>	Nov 21, 2003
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TADALAFIL

<u>AB1</u>	ACCORD HLTHCARE	<u>20MG</u>	<u>A209167 004</u>	Mar 26, 2019
<u>AB1</u>	AJANTA PHARMA LTD	<u>20MG</u>	<u>A209654 004</u>	Mar 26, 2019
<u>AB1</u>	ALEMBIC PHARMS LTD	<u>20MG</u>	<u>A204809 004</u>	Mar 26, 2019
<u>AB1</u>	AMNEAL PHARMS CO	<u>20MG</u>	<u>A209744 004</u>	Mar 26, 2019
<u>AB1</u>	AUROBINDO PHARMA LTD	<u>20MG</u>	<u>A206285 004</u>	Mar 26, 2019
<u>AB1</u>	CIPLA	<u>20MG</u>	<u>A209539 004</u>	Mar 26, 2019
<u>AB1</u>	DR REDDYS LABS LTD	<u>20MG</u>	<u>A210069 004</u>	Mar 26, 2019
<u>AB1</u>	HETERO LABS LTD III	<u>20MG</u>	<u>A209908 004</u>	Mar 26, 2019
<u>AB1</u>	LUPIN LTD	<u>20MG</u>	<u>A210567 004</u>	Mar 26, 2019
<u>AB1</u>	MACLEODS PHARMS LTD	<u>20MG</u>	<u>A207244 004</u>	Oct 07, 2019
<u>AB1</u>	MYLAN	<u>20MG</u>	<u>A206956 003</u>	Apr 29, 2019
<u>AB1</u>	QILU	<u>20MG</u>	<u>A210420 004</u>	Mar 26, 2019
<u>AB1</u>	SUN PHARM	<u>20MG</u>	<u>A208934 004</u>	Mar 26, 2019
<u>AB1</u>	SUNSHINE LAKE	<u>20MG</u>	<u>A211335 004</u>	Mar 26, 2019
<u>AB1</u>	TEVA PHARMS USA	<u>20MG</u>	<u>A090141 004</u>	May 22, 2018
<u>AB1</u>	TORRENT	<u>20MG</u>	<u>A211839 004</u>	Mar 26, 2019
<u>AB1</u>	UNICHEM LABS LTD	<u>20MG</u>	<u>A209250 004</u>	Mar 26, 2019
<u>AB1</u>	ZYDUS PHARMS	<u>20MG</u>	<u>A206693 004</u>	Mar 26, 2019

ADCIRCA

<u>AB2</u>	+!	ELI LILLY CO	<u>20MG</u>	<u>N022332 001</u>	May 22, 2009
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ALYQ

<u>AB2</u>		TEVA PHARMS USA	<u>20MG</u>	<u>A209942 001</u>	Feb 05, 2019
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TADALAFIL

<u>AB2</u>	AJANTA PHARMA LTD	<u>20MG</u>	<u>A210392 001</u>	Feb 05, 2019
<u>AB2</u>	AUROBINDO PHARMA LTD	<u>20MG</u>	<u>A206286 001</u>	Feb 05, 2019
<u>AB2</u>	CIPLA	<u>20MG</u>	<u>A210255 001</u>	Feb 05, 2019
<u>AB2</u>	DR REDDYS LABS LTD	<u>20MG</u>	<u>A210145 001</u>	Feb 05, 2019
<u>AB2</u>	HETERO LABS LTD III	<u>20MG</u>	<u>A209907 001</u>	Feb 05, 2019
<u>AB2</u>	LUPIN LTD	<u>20MG</u>	<u>A210572 001</u>	Feb 05, 2019
<u>AB2</u>	MACLEODS PHARMS LTD	<u>20MG</u>	<u>A207290 001</u>	Oct 16, 2019
<u>AB2</u>	MYLAN	<u>20MG</u>	<u>A200630 001</u>	Aug 03, 2018
<u>AB2</u>	TORRENT	<u>20MG</u>	<u>A212062 001</u>	Mar 26, 2019

PRESCRIPTION DRUG PRODUCT LIST

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** MICRO LABS **0.0015%** **A209051 001** Aug 19, 2019ZIOPTAN**AT** +! OAK PHARMS INC **0.0015%** **N202514 001** Feb 10, 2012TALAZOPARIB TOSYLATE

CAPSULE;ORAL

TALZENNA

+ PFIZER INC EQ 0.25MG BASE N211651 001 Oct 16, 2018

+! EQ 1MG BASE N211651 002 Oct 16, 2018

TALC

AEROSOL;INTRAPLEURAL

SCLEROSOL

+! LYMOL MEDCL 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

TALC

+! LYMOL MEDCL 5GM/BOT N021388 001 Dec 15, 2003

TALIGLUCERASE ALFA

POWDER;INTRAVENOUS

ELELYSO

+! PFIZER 200 UNITS/VIAL N022458 001 May 01, 2012

TAMOXIFEN CITRATE

SOLUTION;ORAL

SOLTAMOX

FORTOVIA

EQ 20MG BASE/10ML N021807 001 Oct 29, 2005

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** MAYNE PHARMA **EQ 10MG BASE** **A075797 001** Feb 20, 2003**AB** ! **EQ 20MG BASE** **A075797 002** Feb 20, 2003**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, 2017TAMSULOSIN HYDROCHLORIDE

CAPSULE;ORAL

FLOMAX**AB** +! SANOFI AVENTIS US **0.4MG** **N020579 001** Apr 15, 1997TAMSULOSIN HYDROCHLORIDE**AB** ALKEM LABS LTD **0.4MG** **A207405 001** Aug 11, 2017**AB** ANBISON LAB **0.4MG** **A211885 001** Oct 17, 2019**AB** ANCHEN PHARMS **0.4MG** **A202010 001** Jan 04, 2013**AB** AUROBINDO PHARMA LTD **0.4MG** **A202433 001** Apr 30, 2013**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

PRESCRIPTION DRUG PRODUCT LISTTAMSULOSIN HYDROCHLORIDE

CAPSULE; ORAL

TAMSULOSIN HYDROCHLORIDE

AB	SUN PHARM INDS LTD	0.4MG	A090931 001	Jul 15, 2010
AB	SYNTHON PHARMS	0.4MG	A078801 001	Apr 27, 2010
AB	TEVA PHARMS	0.4MG	A077630 001	Apr 27, 2010
AB	WOCKHARDT	0.4MG	A078938 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, 2011

TASIMELTEON

CAPSULE; ORAL

HETLIOZ

+	VANDA PHARMS INC	20MG	N205677 001	Jan 31, 2014
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TAVABOROLE

SOLUTION; TOPICAL

KERYDIN

+	ANACOR PHARMS INC	5%	N204427 001	Jul 07, 2014
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TAZAROTENE

AEROSOL, FOAM; TOPICAL

FABIOR

+	MAYNE PHARMA	0.1%	N202428 001	May 11, 2012
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CREAM; TOPICAL

AVAGE

AB	+	ALLERGAN	0.1%	N021184 003	Sep 30, 2002
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TAZAROTENE

AB	ACP NIMBLE	0.1%	A208662 001	Dec 22, 2017
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AB	TARO PHARMS	0.1%	A208258 001	Apr 03, 2017
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TAZORAC

AB	+	ALLERGAN	0.1%	N021184 002	Sep 29, 2000
	+		0.05%	N021184 001	Sep 29, 2000

GEL; TOPICAL

TAZORAC

+	ALLERGAN	0.05%	N020600 001	Jun 13, 1997
+	!	0.1%	N020600 002	Jun 13, 1997

LOTION; TOPICAL

ARAZLO

+	BAUSCH	0.045%	N211882 001	Dec 18, 2019
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TAZEMETOSTAT HYDROBROMIDE

TABLET; ORAL

TAZVERIK

+	EPIZYME INC	EQ 200MG BASE	N211723 001	Jan 23, 2020
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TECHNETIUM TC-99M ALBUMIN AGGREGATED KIT

INJECTABLE; INJECTION

TECHNETIUM TC 99M ALBUMIN AGGREGATED KIT

BS	+	DRAXIMAGE	N/A	N017881 001	Dec 30, 1987
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TECHNETIUM TC-99M BICISATE KIT

INJECTABLE; INJECTION

NEUROLITE

+	LANTHEUS MEDCL	N/A	N020256 001	Nov 23, 1994
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TECHNETIUM TC-99M EXAMETAZIME KIT

INJECTABLE; INJECTION

CERETEC

+	GE HEALTHCARE	N/A	N019829 001	Dec 30, 1988
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PRESCRIPTION DRUG PRODUCT LIST

TECHNETIUM TC-99M EXAMETAZIME KIT

POWDER; INTRAVENOUS

DRAX EXAMETAZIME

JUBILANT DRAXIMAGE N/A

N208870 001 Aug 17, 2017

TECHNETIUM TC-99M MEBROFENIN KIT

INJECTABLE; INJECTION

CHOLETEC**AP** +! BRACCO **N/A****N018963 001** Jan 21, 1987**TECHNETIUM TC-99M MEBROFENIN****AP** PHARMALUCENCE **N/A****A078242 001** Jan 29, 2008TECHNETIUM TC-99M MEDRONATE

INJECTABLE; INJECTION

DRAXIMAGE MDP-25

+! JUBILANT DRAXIMAGE N/A

N018035 002 Feb 27, 2004

TECHNETIUM TC-99M MEDRONATE KIT

INJECTABLE; INJECTION

CIS-MDP

PHARMALUCENCE N/A

N018124 001

TECHNETIUM TC-99M MERTIATIDE KIT

INJECTABLE; INJECTION

TECHNESCAN MAG3**AP** +! CURIUM **N/A****N019882 001** Jun 15, 1990**TECHNETIUM TC99M MERTIATIDE KIT****AP** PHARMALUCENCE **N/A****A208994 001** Jul 12, 2019**AP** SOMMER PHARMS II **N/A****A206489 001** Feb 06, 2020

LLC

TECHNETIUM TC-99M OXIDRONATE KIT

INJECTABLE; INJECTION

TECHNESCAN

+! CURIUM N/A

N018321 001

TECHNETIUM TC-99M PENTETATE KIT

INJECTABLE; INJECTION

DRAXIMAGE DTPA

+! JUBILANT DRAXIMAGE N/A

N018511 001 Dec 29, 1989

TECHNETIUM TC-99M PYROPHOSPHATE KIT

INJECTABLE; INJECTION

CIS-PYRO**AP** PHARMALUCENCE **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** PHARMALUCENCE **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

SOLUTION; INTRAVENOUS, INTRAVESICULAR, OPHTHALMIC

RADIOGENIX SYSTEM

+ NORTHSTAR MEDICAL 30-1153mCi/GENERATOR

N202158 001 Feb 08, 2018

PRESCRIPTION DRUG PRODUCT LISTTECHNETIUM TC-99M SULFUR COLLOID KIT

SOLUTION; INJECTION, ORAL

AN-SULFUR COLLOID

+! PHARMALUCENCE N/A

N017858 001

TECHNETIUM TC-99M TETROFOSMIN KIT

INJECTABLE; INJECTION

MYOVUE 30ML

+! GE HEALTHCARE 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

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 RECOMBINANT

POWDER; SUBCUTANEOUS

GATTEX KIT

+! NPS PHARMS INC 5MG/VIAL

N203441 001 Dec 21, 2012

TEGASEROD MALEATE

TABLET; ORAL

ZELNORM

+! ALFASIGMA EQ 6MG BASE

N021200 002 Jul 24, 2002

TELAVANCIN HYDROCHLORIDE

POWDER; INTRAVENOUS

VIBATIV

+! CUMBERLAND PHARMS EQ 750MG BASE/VIAL

N022110 002 Sep 11, 2009

TELMISARTAN

TABLET; ORAL

MICARDIS**AB** + BOEHRINGER
INGELHEIM **20MG****N020850 003** Apr 04, 2000**AB** + **40MG****N020850 001** Nov 10, 1998**AB** +! **80MG****N020850 002** Nov 10, 1998**TELMISARTAN****AB** ALEMBIC PHARMS LTD **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
LTD **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** 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** MYLAN **20MG****A202397 001** Jul 07, 2014**AB** **40MG****A202397 002** Jul 07, 2014**AB** **80MG****A202397 003** Jul 07, 2014**AB** PRINSTON INC **20MG****A207882 001** May 03, 2017

PRESCRIPTION DRUG PRODUCT LISTTELMISARTAN

TABLET; ORAL

TELMISARTAN

<u>AB</u>		<u>40MG</u>	<u>A207882</u>	<u>002</u>	May 03, 2017
<u>AB</u>		<u>80MG</u>	<u>A207882</u>	<u>003</u>	May 03, 2017
<u>AB</u>	SANDOZ INC	<u>20MG</u>	<u>A203867</u>	<u>001</u>	Nov 03, 2014
<u>AB</u>		<u>40MG</u>	<u>A203867</u>	<u>002</u>	Nov 03, 2014
<u>AB</u>		<u>80MG</u>	<u>A203867</u>	<u>003</u>	Nov 03, 2014
<u>AB</u>	TORRENT	<u>20MG</u>	<u>A203171</u>	<u>001</u>	Jul 07, 2014
<u>AB</u>		<u>40MG</u>	<u>A203171</u>	<u>002</u>	Jul 07, 2014
<u>AB</u>		<u>80MG</u>	<u>A203171</u>	<u>003</u>	Jul 07, 2014
<u>AB</u>	ZYDUS PHARMS	<u>20MG</u>	<u>A203325</u>	<u>001</u>	Aug 26, 2014
<u>AB</u>		<u>40MG</u>	<u>A203325</u>	<u>002</u>	Aug 26, 2014
<u>AB</u>		<u>80MG</u>	<u>A203325</u>	<u>003</u>	Aug 26, 2014

TELOTRISTAT ETIPRATE

TABLET; ORAL

XERMELO

+! LEXICON PHARMS INC EQ 250MG BASE N208794 001 Feb 28, 2017

TEMAZEPAM

CAPSULE; ORAL

RESTORIL

<u>AB</u>	+	SPECGX LLC	<u>7.5MG</u>	<u>N018163</u>	<u>003</u>	Oct 25, 1991
<u>AB</u>	+		<u>15MG</u>	<u>N018163</u>	<u>001</u>	
<u>AB</u>	+		<u>22.5MG</u>	<u>N018163</u>	<u>004</u>	Nov 02, 2004
<u>AB</u>	+!		<u>30MG</u>	<u>N018163</u>	<u>002</u>	

TEMAZEPAM

<u>AB</u>		ACTAVIS ELIZABETH	<u>15MG</u>	<u>A071620</u>	<u>002</u>	Aug 07, 1987
<u>AB</u>			<u>30MG</u>	<u>A071620</u>	<u>001</u>	Aug 07, 1987
<u>AB</u>		ALEMBIC PHARMS LTD	<u>7.5MG</u>	<u>A211542</u>	<u>001</u>	Nov 23, 2018
<u>AB</u>			<u>15MG</u>	<u>A211542</u>	<u>002</u>	Nov 23, 2018
<u>AB</u>			<u>22.5MG</u>	<u>A211542</u>	<u>003</u>	Nov 23, 2018
<u>AB</u>			<u>30MG</u>	<u>A211542</u>	<u>004</u>	Nov 23, 2018
<u>AB</u>		AMNEAL PHARMS	<u>7.5MG</u>	<u>A203482</u>	<u>001</u>	May 23, 2016
<u>AB</u>			<u>15MG</u>	<u>A203482</u>	<u>002</u>	May 23, 2016
<u>AB</u>			<u>22.5MG</u>	<u>A203482</u>	<u>003</u>	May 23, 2016
<u>AB</u>			<u>30MG</u>	<u>A203482</u>	<u>004</u>	May 23, 2016
<u>AB</u>		MYLAN	<u>7.5MG</u>	<u>A070920</u>	<u>002</u>	May 21, 2010
<u>AB</u>			<u>15MG</u>	<u>A070920</u>	<u>004</u>	Jul 07, 1986
<u>AB</u>			<u>22.5MG</u>	<u>A070920</u>	<u>003</u>	Jun 12, 2009
<u>AB</u>			<u>30MG</u>	<u>A070920</u>	<u>001</u>	Jul 10, 1986
<u>AB</u>		NOVEL LABS INC	<u>7.5MG</u>	<u>A071457</u>	<u>002</u>	Jun 22, 2012
<u>AB</u>			<u>15MG</u>	<u>A071456</u>	<u>001</u>	Apr 21, 1987
<u>AB</u>			<u>22.5MG</u>	<u>A071457</u>	<u>003</u>	Jun 22, 2012
<u>AB</u>			<u>30MG</u>	<u>A071457</u>	<u>001</u>	Apr 21, 1987
<u>AB</u>		PRINSTON INC	<u>7.5MG</u>	<u>A201781</u>	<u>001</u>	Jun 04, 2015
<u>AB</u>			<u>15MG</u>	<u>A201781</u>	<u>002</u>	Jun 04, 2015
<u>AB</u>			<u>22.5MG</u>	<u>A201781</u>	<u>003</u>	Jun 04, 2015
<u>AB</u>			<u>30MG</u>	<u>A201781</u>	<u>004</u>	Jun 04, 2015
<u>AB</u>		SANDOZ	<u>15MG</u>	<u>A071427</u>	<u>001</u>	Jan 12, 1988
<u>AB</u>			<u>30MG</u>	<u>A071428</u>	<u>001</u>	Jan 12, 1988
<u>AB</u>		SUN PHARM INDUSTRIES	<u>7.5MG</u>	<u>A078581</u>	<u>001</u>	Sep 08, 2009
<u>AB</u>			<u>22.5MG</u>	<u>A071175</u>	<u>002</u>	Sep 14, 2009

TEMOZOLOMIDE

CAPSULE; ORAL

TEMODAR

<u>AB</u>	+	MERCK SHARP DOHME	<u>5MG</u>	<u>N021029</u>	<u>001</u>	Aug 11, 1999
<u>AB</u>	+		<u>20MG</u>	<u>N021029</u>	<u>002</u>	Aug 11, 1999
<u>AB</u>	+		<u>100MG</u>	<u>N021029</u>	<u>003</u>	Aug 11, 1999
<u>AB</u>	+		<u>140MG</u>	<u>N021029</u>	<u>005</u>	Oct 19, 2006
<u>AB</u>	+		<u>180MG</u>	<u>N021029</u>	<u>006</u>	Oct 19, 2006
<u>AB</u>	+!		<u>250MG</u>	<u>N021029</u>	<u>004</u>	Aug 11, 1999

TEMOZOLOMIDE

<u>AB</u>		ACCORD HLTHCARE	<u>5MG</u>	<u>A201528</u>	<u>001</u>	Feb 27, 2017
<u>AB</u>			<u>20MG</u>	<u>A201528</u>	<u>002</u>	Feb 27, 2017
<u>AB</u>			<u>100MG</u>	<u>A201528</u>	<u>003</u>	Feb 27, 2017
<u>AB</u>			<u>140MG</u>	<u>A201528</u>	<u>004</u>	Feb 27, 2017
<u>AB</u>			<u>180MG</u>	<u>A201528</u>	<u>005</u>	Feb 27, 2017
<u>AB</u>			<u>250MG</u>	<u>A201528</u>	<u>006</u>	Feb 27, 2017
<u>AB</u>		AMERIGEN PHARMS LTD	<u>5MG</u>	<u>A203490</u>	<u>001</u>	Jul 13, 2016
<u>AB</u>			<u>20MG</u>	<u>A203490</u>	<u>002</u>	Jul 13, 2016

PRESCRIPTION DRUG PRODUCT LIST

TEMOZOLOMIDE

CAPSULE; ORAL

TEMOZOLOMIDE

<u>AB</u>		<u>100MG</u>	<u>A203490 003</u>	Jul 13, 2016
<u>AB</u>		<u>140MG</u>	<u>A203490 004</u>	Jul 13, 2016
<u>AB</u>		<u>180MG</u>	<u>A203490 005</u>	Jul 13, 2016
<u>AB</u>		<u>250MG</u>	<u>A203490 006</u>	Jul 13, 2016
<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>	BARR	<u>5MG</u>	<u>A078879 001</u>	Mar 01, 2010
<u>AB</u>		<u>20MG</u>	<u>A078879 002</u>	Mar 01, 2010
<u>AB</u>		<u>100MG</u>	<u>A078879 003</u>	Mar 01, 2010
<u>AB</u>		<u>140MG</u>	<u>A078879 005</u>	Mar 01, 2010
<u>AB</u>		<u>180MG</u>	<u>A078879 006</u>	Mar 01, 2010
<u>AB</u>		<u>250MG</u>	<u>A078879 004</u>	Mar 01, 2010
<u>AB</u>	CHEMI SPA	<u>5MG</u>	<u>A204639 001</u>	Nov 23, 2016
<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>	IDT AUSTRALIA LTD	<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>	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>	WATSON LABS TEVA	<u>5MG</u>	<u>A203959 001</u>	Apr 18, 2017
<u>AB</u>		<u>20MG</u>	<u>A203959 002</u>	Apr 18, 2017
<u>AB</u>		<u>100MG</u>	<u>A203959 003</u>	Apr 18, 2017
<u>AB</u>		<u>140MG</u>	<u>A203959 004</u>	Apr 18, 2017
<u>AB</u>		<u>250MG</u>	<u>A203959 005</u>	Apr 18, 2017
<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>TORISEL</u>				
<u>AP</u>	+! PF PRISM CV	<u>25MG/ML (25MG/ML)</u>	<u>N022088 001</u>	May 30, 2007

PRESCRIPTION DRUG PRODUCT LIST

TENAPANOR HYDROCHLORIDE

TABLET;ORAL

IBSRELA

+! ARDELYX INC EQ 50MG BASE N211801 001 Sep 12, 2019

TENOFOVIR ALAFENAMIDE FUMARATE

TABLET;ORAL

VEMLIDY

+! GILEAD SCIENCES INC EQ 25MG BASE N208464 001 Nov 10, 2016

TENOFOVIR DISOPROXIL FUMARATE

POWDER;ORAL

VIREAD

+! GILEAD SCIENCES INC 40MG/SCOOPFUL N022577 001 Jan 18, 2012

TABLET;ORAL

TENOFOVIR DISOPROXIL FUMARATE

AB	AUROBINDO PHARMA LTD	150MG	A090647 001	Jan 26, 2018
AB		200MG	A090647 002	Jan 26, 2018
AB		250MG	A090647 003	Jan 26, 2018
AB		300MG	A090647 004	Jan 26, 2018
AB	CASI PHARMS INC	300MG	A209550 001	Feb 26, 2018
AB	CIPLA	300MG	A078800 001	Jan 26, 2018
AB	HETERO LABS LTD III	300MG	A090636 001	Jan 26, 2018
AB	MACLEODS PHARMS LTD	300MG	A203232 001	Jan 26, 2018
AB	QILU	200MG	A209498 001	Mar 02, 2018
AB		250MG	A209498 002	Mar 02, 2018
AB		300MG	A209498 003	Mar 02, 2018
AB	STRIDES PHARMA	300MG	A090742 001	Jan 26, 2018
AB	TEVA PHARMS USA	150MG	A091612 002	Jan 26, 2018
AB		200MG	A091612 003	Jan 26, 2018
AB		250MG	A091612 004	Jan 26, 2018
AB		300MG	A091612 001	Mar 18, 2015
VIREAD				
AB	+ GILEAD SCIENCES INC	150MG	N021356 002	Jan 18, 2012
AB	+	200MG	N021356 003	Jan 18, 2012
AB	+	250MG	N021356 004	Jan 18, 2012
AB	+!	300MG	N021356 001	Oct 26, 2001

TERAZOSIN HYDROCHLORIDE

CAPSULE;ORAL

TERAZOSIN HYDROCHLORIDE

AB	APNAR PHARMA LP	EQ 1MG BASE	A074823 001	Mar 30, 1998
AB	!	EQ 2MG BASE	A074823 002	Mar 30, 1998
AB		EQ 5MG BASE	A074823 003	Mar 30, 1998
AB		EQ 10MG BASE	A074823 004	Mar 30, 1998
AB	HIKMA	EQ 1MG BASE	A075498 001	Apr 12, 2001
AB		EQ 2MG BASE	A075498 002	Apr 12, 2001
AB		EQ 5MG BASE	A075498 003	Apr 12, 2001
AB		EQ 10MG BASE	A075498 004	Apr 12, 2001
AB	JUBILANT CADISTA	EQ 1MG BASE	A075317 001	Dec 20, 2004
AB		EQ 2MG BASE	A075317 002	Dec 20, 2004
AB		EQ 5MG BASE	A075317 003	Dec 20, 2004
AB		EQ 10MG BASE	A075317 004	Dec 20, 2004
AB	MYLAN	EQ 1MG BASE	A075140 002	Feb 11, 2000
AB		EQ 2MG BASE	A075140 003	Feb 11, 2000
AB		EQ 5MG BASE	A075140 001	Feb 11, 2000
AB		EQ 10MG BASE	A075140 004	Feb 11, 2000

TERBINAFINE HYDROCHLORIDE

TABLET;ORAL

LAMISIL

AB	+! NOVARTIS	EQ 250MG BASE	N020539 001	May 10, 1996
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TERBINAFINE HYDROCHLORIDE

AB	AUROBINDO PHARMA	EQ 250MG BASE	A078297 001	Jul 02, 2007
AB	BRECKENRIDGE PHARM	EQ 250MG BASE	A077714 001	Jun 04, 2010
AB	CIPLA	EQ 250MG BASE	A077137 001	Jul 02, 2007
AB	DR REDDYS LABS INC	EQ 250MG BASE	A076390 001	Jul 02, 2007
AB	GLENMARK GENERICS	EQ 250MG BASE	A078157 001	Jul 02, 2007
AB	HARRIS PHARM	EQ 250MG BASE	A077919 001	Jul 02, 2007
AB	HERITAGE PHARMA	EQ 250MG BASE	A076377 001	Jul 02, 2007
AB	INVAGEN PHARMS	EQ 250MG BASE	A077533 001	Jul 02, 2007
AB	ORCHID HLTHCARE	EQ 250MG BASE	A078163 001	Jul 02, 2007

PRESCRIPTION DRUG PRODUCT LIST

TERBUTALINE SULFATE

INJECTABLE; INJECTION

TERBUTALINE SULFATE

AP	AKORN	<u>1MG/ML</u>	<u>A078151</u>	<u>001</u>	Jan 07, 2008
AP	! ATHENEX INC	<u>1MG/ML</u>	<u>A076770</u>	<u>001</u>	Apr 23, 2004
AP	FRESENIUS KABI USA	<u>1MG/ML</u>	<u>A076887</u>	<u>001</u>	May 26, 2004
AP	HIKMA FARMACEUTICA	<u>1MG/ML</u>	<u>A078630</u>	<u>001</u>	May 20, 2009
AP	UNITED BIOMEDCL	<u>1MG/ML</u>	<u>A200122</u>	<u>001</u>	Nov 08, 2013

TABLET; ORAL

BRETHINE

AB	+ ANI PHARMS INC	<u>2.5MG</u>	<u>N017849</u>	<u>001</u>	
AB	+	<u>5MG</u>	<u>N017849</u>	<u>002</u>	
<u>TERBUTALINE SULFATE</u>					
AB	IMPAX LABS	<u>2.5MG</u>	<u>A075877</u>	<u>001</u>	Jun 26, 2001
AB		<u>5MG</u>	<u>A075877</u>	<u>002</u>	Jun 26, 2001
AB	LANNETT CO INC	<u>2.5MG</u>	<u>A077152</u>	<u>001</u>	Mar 25, 2005
AB	!	<u>5MG</u>	<u>A077152</u>	<u>002</u>	Mar 25, 2005

TERCONAZOLE

CREAM; VAGINAL

TERCONAZOLE

AB	FOUGERA PHARMS	<u>0.4%</u>	<u>A076712</u>	<u>001</u>	Feb 18, 2005
AB	! TARO	<u>0.4%</u>	<u>A076043</u>	<u>001</u>	Jan 19, 2005
BX	+! FOUGERA PHARMS INC	0.8%	N021735	001	Oct 01, 2004
BX	! TARO	0.8%	A075953	001	Apr 06, 2004

SUPPOSITORY; VAGINAL

TERCONAZOLE

AB	! PERRIGO ISRAEL	<u>80MG</u>	<u>A077149</u>	<u>001</u>	Mar 17, 2006
AB	TARO	<u>80MG</u>	<u>A077553</u>	<u>001</u>	Mar 09, 2007

TERIFLUNOMIDE

TABLET; ORAL

AUBAGIO

AB	+ SANOFI AVENTIS US	<u>7MG</u>	<u>N202992</u>	<u>001</u>	Sep 12, 2012
AB	+!	<u>14MG</u>	<u>N202992</u>	<u>002</u>	Sep 12, 2012

TERIFLUNOMIDE

AB	ACCORD HLTHCARE	<u>7MG</u>	<u>A209690</u>	<u>001</u>	Jan 07, 2019
AB		<u>14MG</u>	<u>A209690</u>	<u>002</u>	Jan 07, 2019
AB	ALEMBIC PHARMS LTD	<u>7MG</u>	<u>A209572</u>	<u>001</u>	Apr 19, 2019
AB		<u>14MG</u>	<u>A209572</u>	<u>002</u>	Apr 19, 2019
AB	GLENMARK PHARMS	<u>7MG</u>	<u>A209663</u>	<u>001</u>	Nov 15, 2018
AB		<u>14MG</u>	<u>A209663</u>	<u>002</u>	Nov 15, 2018
AB	MYLAN	<u>7MG</u>	<u>A209702</u>	<u>001</u>	Feb 28, 2020
AB		<u>14MG</u>	<u>A209702</u>	<u>002</u>	Feb 28, 2020
AB	SANDOZ INC	<u>7MG</u>	<u>A209710</u>	<u>001</u>	Jan 03, 2019
AB		<u>14MG</u>	<u>A209710</u>	<u>002</u>	Jan 03, 2019

TERIPARATIDE

SOLUTION; SUBCUTANEOUS

BONSITY

ALVOGEN

0.62MG/2.48ML (0.25MG/ML)

N211939 001 Oct 04, 2019

FORTEO

+! LILLY

0.6MG/2.4ML (0.25MG/ML)

N021318 002 Jun 25, 2008

TESAMORELIN ACETATE

POWDER; SUBCUTANEOUS

EGRIFTA

+! THERATECHNOLOGIES

EQ 1MG BASE/VIAL

N022505 001 Nov 10, 2010

+!

EQ 2MG BASE/VIAL

N022505 002 Nov 29, 2011

TESTOSTERONE

FILM, EXTENDED RELEASE; TRANSDERMAL

ANDRODERM

+! ALLERGAN

2MG/24HR

N020489 003 Oct 20, 2011

+!

4MG/24HR

N020489 004 Oct 20, 2011

GEL; TRANSDERMAL

ANDROGEL

AB1	+ ABBVIE	<u>25MG/2.5GM PACKET</u>	<u>N021015</u>	<u>001</u>	Feb 28, 2000
AB1	+!	<u>50MG/5GM PACKET</u>	<u>N021015</u>	<u>002</u>	Feb 28, 2000

TESTOSTERONE

AB1	ACTAVIS LABS UT INC	<u>25MG/2.5GM PACKET</u>	<u>A076737</u>	<u>001</u>	Jan 27, 2006
AB1		<u>50MG/5GM PACKET</u>	<u>A076737</u>	<u>002</u>	Jan 27, 2006
AB1	PAR PHARM	<u>50MG/5GM PACKET</u>	<u>A076744</u>	<u>002</u>	May 23, 2007

PRESCRIPTION DRUG PRODUCT LIST

TESTOSTERONE

GEL; TRANSDERMAL

ANDROGEL

AB2 + ABBVIE 1.62% (20.25MG/1.25GM PACKET) **N022309 002** Sep 07, 2012
AB2 +! 1.62% (40.5MG/2.5GM PACKET) **N022309 003** Sep 07, 2012

TESTIM

AB2 +! AUXILIUM PHARMS LLC 50MG/5GM PACKET **N021454 001** Oct 31, 2002

TESTOSTERONE

AB2 ACTAVIS LABS UT INC 50MG/5GM PACKET **A091073 001** Sep 18, 2017
AB2 PERRIGO UK FINCO 1.62% (20.25MG/1.25GM PACKET) **A205781 001** Jul 12, 2017
AB2 1.62% (40.5MG/2.5GM PACKET) **A205781 002** Jul 12, 2017

VOGELXO

AB2 UPSHER SMITH LABS 50MG/5GM PACKET **N204399 002** Jun 04, 2014

GEL, METERED; NASAL

NATESTO

ACERUS 5.5MG/0.122GM ACTUATION N205488 001 May 28, 2014

GEL, METERED; TRANSDERMAL

ANDROGEL

AB +! ABBVIE 1.62% (20.25MG/1.25GM ACTUATION) **N022309 001** Apr 29, 2011
AB +! 12.5MG/1.25GM ACTUATION **N021015 003** Sep 26, 2003

FORTESTA

AB +! ENDO PHARMS 10MG/0.5GM ACTUATION **N021463 001** Dec 29, 2010

TESTOSTERONE

AB ACTAVIS LABS UT INC 1.62% (20.25MG/1.25GM ACTUATION) **A204570 001** Apr 10, 2019
AB 10MG/0.5GM ACTUATION **A204571 001** Aug 05, 2015
AB 12.5MG/1.25GM ACTUATION **A076737 003** Mar 09, 2015
AB AMNEAL PHARMS LLC 1.62% (20.25MG/1.25GM ACTUATION) **A207373 001** Apr 10, 2019
AB DR REDDYS 1.62% (20.25MG/1.25GM ACTUATION) **A208620 001** Apr 10, 2019
AB LUPIN ATLANTIS 1.62% (20.25MG/1.25GM ACTUATION) **A208560 001** Apr 10, 2019
AB PERRIGO ISRAEL 1.62% (20.25MG/1.25GM ACTUATION) **A204268 001** Aug 04, 2015
AB TWI PHARMS 1.62% (20.25MG/1.25GM ACTUATION) **A209390 001** Sep 23, 2019

VOGELXO

BX UPSHER SMITH LABS 12.5MG/1.25GM ACTUATION N204399 003 Jun 04, 2014

PELLET; IMPLANTATION

TESTOPEL

! AUXILIUM PHARMS INC 75MG A080911 001

SOLUTION, METERED; TRANSDERMAL

TESTOSTERONE

AT ACTAVIS LABS UT INC 30MG/1.5ML ACTUATION **A205328 001** Aug 07, 2017
AT CIPLA 30MG/1.5ML ACTUATION **A209533 001** Jan 29, 2018
AT LUPIN LTD 30MG/1.5ML ACTUATION **A208061 001** Oct 23, 2017
AT ! PERRIGO ISRAEL 30MG/1.5ML ACTUATION **A204255 001** Feb 28, 2017

TESTOSTERONE CYPIONATE

INJECTABLE; INJECTION

DEPO-TESTOSTERONE

AO ! PHARMACIA AND 100MG/ML **A085635 002**
 UPJOHN

AO ! 200MG/ML **A085635 003**

TESTOSTERONE CYPIONATE

AO AM REGENT 200MG/ML **A207742 001** Jun 16, 2017
AO CIPLA 100MG/ML **A210362 001** Jun 19, 2018
AO 200MG/ML **A210362 002** Jun 19, 2018
AO HIKMA FARMACEUTICA 200MG/ML **A091244 001** May 01, 2012
AO PADDOCK LLC 200MG/ML **A040530 001** Jan 31, 2005
AO SANDOZ INC 100MG/ML **A040615 001** Aug 10, 2006
AO 200MG/ML **A040615 002** Aug 10, 2006
AO SUN PHARM INDS LTD 100MG/ML **A201720 001** Jun 03, 2013
AO 200MG/ML **A201720 002** Jun 03, 2013
AO WATSON PHARMS INC 200MG/ML **A086030 001**
AO WEST-WARD PHARMS 100MG/ML **A090387 001** Jul 15, 2010
AO INT 200MG/ML **A090387 002** Jul 15, 2010
AO WILSHIRE PHARMS INC 200MG/ML **A206368 001** Apr 24, 2019

TESTOSTERONE ENANTHATE

INJECTABLE; INJECTION

TESTOSTERONE ENANTHATE

AO HIKMA FARMACEUTICA 200MG/ML **A091120 001** Sep 18, 2012
AO NEXUS PHARMS 200MG/ML **A040575 001** Jun 14, 2006
AO ! WATSON PHARMS INC 200MG/ML **A085598 001**

PRESCRIPTION DRUG PRODUCT LISTTESTOSTERONE ENANTHATE

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

+		CLARUS	158MG	N206089	001	Mar 27, 2019
+			198MG	N206089	002	Mar 27, 2019
+	!		237MG	N206089	003	Mar 27, 2019

INJECTABLE;INTRAMUSCULAR

AVEED

+	!	ENDO PHARMS INC	750MG/3ML (250MG/ML)	N022219	001	Mar 05, 2014
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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>		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 INC	<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>		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>		HIKMA	<u>12.5MG</u>	<u>A209739</u>	<u>001</u>	Apr 08, 2019
<u>AB</u>			<u>25MG</u>	<u>A209739</u>	<u>002</u>	Apr 08, 2019
<u>AB</u>		LUPIN LTD	<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>		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
<u>AB</u>		PIRAMAL HLTHCARE UK	<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>		SUN PHARM	<u>12.5MG</u>	<u>A206129</u>	<u>001</u>	Aug 17, 2015
<u>AB</u>			<u>25MG</u>	<u>A206129</u>	<u>002</u>	Aug 17, 2015

XENAZINE

<u>AB</u>	+	VALEANT PHARMS NORTH	<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

+	!	ALCON LABS	0.5%	N208135	001	Feb 29, 2016
		BAUSCH	0.5%	N210821	001	Mar 12, 2019

TETRACYCLINE HYDROCHLORIDE

CAPSULE;ORAL

ACHROMYCIN V

<u>AB</u>	+	AVET	<u>250MG</u>	<u>N050278</u>	<u>003</u>	
<u>AB</u>	+	!	<u>500MG</u>	<u>N050278</u>	<u>001</u>	

TETRACYCLINE HYDROCHLORIDE

<u>AB</u>		AMNEAL PHARMS NY	<u>250MG</u>	<u>A210674</u>	<u>001</u>	Sep 18, 2018
<u>AB</u>			<u>500MG</u>	<u>A210674</u>	<u>002</u>	Sep 18, 2018
<u>AB</u>		BRECKENRIDGE	<u>250MG</u>	<u>A210662</u>	<u>001</u>	Nov 07, 2018
<u>AB</u>			<u>500MG</u>	<u>A210662</u>	<u>002</u>	Nov 07, 2018
<u>AB</u>		CHARTWELL TETRA	<u>250MG</u>	<u>A062752</u>	<u>001</u>	Aug 12, 1988
<u>AB</u>			<u>500MG</u>	<u>A062752</u>	<u>002</u>	Aug 12, 1988
<u>AB</u>		STRIDES PHARMA	<u>250MG</u>	<u>A212635</u>	<u>001</u>	Mar 03, 2020
<u>AB</u>			<u>500MG</u>	<u>A212635</u>	<u>002</u>	Mar 03, 2020
<u>AB</u>		WATSON LABS	<u>250MG</u>	<u>A061837</u>	<u>001</u>	
<u>AB</u>			<u>500MG</u>	<u>A061837</u>	<u>002</u>	

TETRAHYDROZOLINE HYDROCHLORIDE

SOLUTION;NASAL

TYZINE

!		FOUGERA PHARMS	0.05%	A086576	002	
			0.1%	A086576	001	

SPRAY;NASAL

TYZINE

!		FOUGERA PHARMS	0.1%	A086576	003	
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PRESCRIPTION DRUG PRODUCT LIST

3-421 (of 453)

THALIDOMIDE

CAPSULE; ORAL

THALOMID

+	CELGENE	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

AP	+	CURIUM	<u>1mCi/ML</u>	<u>N018150 001</u>	
AP	+	GE HEALTHCARE	<u>1mCi/ML</u>	<u>N018110 002</u>	Feb 27, 1996
AP	+	LANTHEUS MEDCL	<u>1mCi/ML</u>	<u>N017806 001</u>	

THEOPHYLLINE

CAPSULE, EXTENDED RELEASE; ORAL

THEO-24

		ACTIENT PHARMS	100MG	A087942 001	Aug 22, 1983
!		AUXILIUM PHARMS INC	400MG	A081034 001	Feb 28, 1992
		AUXILIUM PHARMS LLC	200MG	A087943 001	Aug 22, 1983
			300MG	A087944 001	Aug 22, 1983

INJECTABLE; INJECTION

THEOPHYLLINE 0.04% AND DEXTROSE 5% IN PLASTIC CONTAINER

+	B BRAUN	40MG/100ML	N019826 001	Aug 14, 1992
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THEOPHYLLINE 0.08% AND DEXTROSE 5% IN PLASTIC CONTAINER

+	B BRAUN	80MG/100ML	N019826 002	Aug 14, 1992
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THEOPHYLLINE 0.16% AND DEXTROSE 5% IN PLASTIC CONTAINER

+	B BRAUN	160MG/100ML	N019826 003	Aug 14, 1992
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THEOPHYLLINE 0.32% AND DEXTROSE 5% IN PLASTIC CONTAINER

+	B BRAUN	320MG/100ML	N019826 006	Aug 14, 1992
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SOLUTION; ORAL

THEOPHYLLINE

AA	!	LANNETT CO INC	<u>80MG/15ML</u>	<u>A091156 001</u>	Apr 13, 2011
AA		TRIS PHARMA INC	<u>80MG/15ML</u>	<u>A091586 001</u>	Jun 15, 2012

SOLUTION, ELIXIR; ORAL

ELIXOPHYLLIN

AA	!	NOSTRUM LABS INC	<u>80MG/15ML</u>	<u>A085186 001</u>	
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THEOPHYLLINE

AA		PHARM ASSOC	<u>80MG/15ML</u>	<u>A206344 001</u>	Dec 16, 2016
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TABLET, EXTENDED RELEASE; ORAL

THEOCHRON

AB		NOSTRUM PHARMS LLC	<u>100MG</u>	<u>A087400 003</u>	Feb 21, 1985
AB			<u>200MG</u>	<u>A087400 004</u>	Feb 21, 1985

THEOPHYLLINE

AB		ALEMBIC PHARMS LTD	<u>300MG</u>	<u>A090430 001</u>	Oct 27, 2010
AB		GLENMARK GENERICS	<u>400MG</u>	<u>A090355 001</u>	Jul 13, 2010
AB			<u>600MG</u>	<u>A090355 002</u>	Jul 13, 2010
AB	!	HERITAGE PHARMA	<u>100MG</u>	<u>A089807 001</u>	Apr 30, 1990
AB	!		<u>200MG</u>	<u>A089808 001</u>	Apr 30, 1990
AB			<u>300MG</u>	<u>A089763 001</u>	Apr 30, 1990
AB		NOSTRUM LABS INC	<u>400MG</u>	<u>A040560 003</u>	Apr 21, 2006
AB	!		<u>600MG</u>	<u>A040560 002</u>	Apr 21, 2006
AB		RHODES PHARMS	<u>400MG</u>	<u>A040086 002</u>	Sep 01, 1982
AB			<u>600MG</u>	<u>A040086 001</u>	Apr 15, 1996
	!	ALEMBIC PHARMS LTD	450MG	A090430 002	Oct 27, 2010

THIAMINE HYDROCHLORIDE

INJECTABLE; INJECTION

THIAMINE HYDROCHLORIDE

AP	!	FRESENIUS KABI USA	<u>100MG/ML</u>	<u>A080556 001</u>	
AP		MYLAN INSTITUTIONAL	<u>100MG/ML</u>	<u>A091623 001</u>	Jun 25, 2012
AP		SAGENT PHARMS INC	<u>100MG/ML</u>	<u>A206106 001</u>	Dec 01, 2017

THIOGUANINE

TABLET; ORAL

THIOGUANINE

+	ASPEN GLOBAL INC	40MG	N012429 001	
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PRESCRIPTION DRUG PRODUCT LIST

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>	DR REDDYS	<u>15MG/VIAL</u>	<u>A210337 001</u>	May 04, 2018
<u>AP</u>	JIANGSU HENGRUI MED	<u>15MG/VIAL</u>	<u>A209150 001</u>	May 04, 2018
<u>AP</u>	STI PHARMA LLC	<u>15MG/VIAL</u>	<u>A208242 001</u>	Jan 10, 2020
<u>AP</u>	! WEST-WARD PHARMS INT	<u>15MG/VIAL</u>	<u>A075547 001</u>	Apr 02, 2001

POWDER; INTRAVENOUS

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>	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

THIOTHIXENE

CAPSULE; ORAL

THIOTHIXENE

<u>AB</u>	MYLAN	<u>1MG</u>	<u>A071093 002</u>	Jun 23, 1987
<u>AB</u>		<u>2MG</u>	<u>A071093 003</u>	Jun 23, 1987
<u>AB</u>		<u>5MG</u>	<u>A071093 004</u>	Jun 23, 1987
<u>AB</u>		<u>10MG</u>	<u>A071093 001</u>	Jun 23, 1987
<u>AB</u>	NOVITIUM PHARMA	<u>1MG</u>	<u>A211642 001</u>	Apr 05, 2019
<u>AB</u>		<u>2MG</u>	<u>A211642 002</u>	Apr 05, 2019
<u>AB</u>	!	<u>5MG</u>	<u>A211642 003</u>	Apr 05, 2019
<u>AB</u>		<u>10MG</u>	<u>A211642 004</u>	Apr 05, 2019

THYROTROPIN ALFA

INJECTABLE; INJECTION

THYROGEN

+!	GENZYME	1.1MG/VIAL	N020898 001	Nov 30, 1998
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TIAGABINE HYDROCHLORIDE

TABLET; ORAL

GABITRIL

<u>AB</u>	+ CEPHALON	<u>2MG</u>	<u>N020646 005</u>	Apr 16, 1999
<u>AB</u>	+!	<u>4MG</u>	<u>N020646 001</u>	Sep 30, 1997
<u>AB</u>	+	<u>12MG</u>	<u>N020646 002</u>	Sep 30, 1997
<u>AB</u>	+	<u>16MG</u>	<u>N020646 003</u>	Sep 30, 1997

TIAGABINE HYDROCHLORIDE

<u>AB</u>	AMNEAL PHARMS CO	<u>2MG</u>	<u>A208181 001</u>	Dec 08, 2017
<u>AB</u>		<u>4MG</u>	<u>A208181 002</u>	Dec 08, 2017
<u>AB</u>		<u>12MG</u>	<u>A208181 003</u>	Dec 08, 2017
<u>AB</u>		<u>16MG</u>	<u>A208181 004</u>	Dec 08, 2017
<u>AB</u>	SUN PHARM INDS	<u>2MG</u>	<u>A077555 001</u>	Nov 04, 2011
<u>AB</u>		<u>4MG</u>	<u>A077555 002</u>	Nov 04, 2011
<u>AB</u>	WILSHIRE PHARMS INC	<u>2MG</u>	<u>A206857 001</u>	Oct 13, 2017
<u>AB</u>		<u>4MG</u>	<u>A206857 002</u>	Oct 13, 2017
<u>AB</u>		<u>12MG</u>	<u>A206857 003</u>	Oct 13, 2017
<u>AB</u>		<u>16MG</u>	<u>A206857 004</u>	Oct 13, 2017

TICAGRELOR

TABLET; ORAL

BRILINTA

<u>AB</u>	+! ASTRAZENECA PHARMS	<u>90MG</u>	<u>N022433 001</u>	Jul 20, 2011
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TICAGRELOR

<u>AB</u>	HISUN PHARM HANGZHOU	<u>90MG</u>	<u>A208575 001</u>	Jan 23, 2019
	BRILINTA			
	+ ASTRAZENECA PHARMS	60MG	N022433 002	Sep 03, 2015

PRESCRIPTION DRUG PRODUCT LIST

TICLOPIDINE HYDROCHLORIDE

TABLET; ORAL

TICLOPIDINE HYDROCHLORIDE

AB	APOTEX	250MG	A075089 001	Jul 01, 1999
AB	! TEVA	250MG	A075149 001	Aug 20, 1999

TIGECYCLINE

POWDER; INTRAVENOUS

TIGECYCLINE

AP	AMNEAL PHARMS LLC	50MG/VIAL	N211158 001	Aug 02, 2018
AP	APOTEX	50MG/VIAL	A204439 001	Dec 21, 2018
AP	AUROBINDO PHARMA LTD	50MG/VIAL	A206335 001	Jun 11, 2019
AP	FRESENIUS KABI USA	50MG/VIAL	N205645 001	Dec 01, 2016
AP	SANDOZ INC	50MG/VIAL	A091620 001	May 27, 2015
AP	XELLIA PHARMS APS	50MG/VIAL	A205722 001	Oct 18, 2019

TYGACIL

AP	! PF PRISM CV	50MG/VIAL	N021821 001	Jun 15, 2005
<u>TIGECYCLINE</u>				
	ACCORD HLTHCARE INC	50MG/VIAL	N208744 001	Jan 18, 2018

TIMOLOL

SOLUTION/DROPS; OPHTHALMIC

BETIMOL

AT	! OAK PHARMS INC	EQ 0.25% BASE	N020439 001	Mar 31, 1995
AT	!	EQ 0.5% BASE	N020439 002	Mar 31, 1995

TIMOLOL

AT	AKORN	EQ 0.25% BASE	A205309 001	Sep 30, 2016
AT		EQ 0.5% BASE	A205309 002	Sep 30, 2016

TIMOLOL MALEATE

SOLUTION, GEL FORMING/DROPS; OPHTHALMIC

TIMOLOL MALEATE

AB	! SANDOZ INC	EQ 0.25% BASE	N020963 001	Oct 21, 1998
AB	!	EQ 0.5% BASE	N020963 002	Oct 21, 1998

TIMOPTIC-XE

AB	! VALEANT PHARMS LLC	EQ 0.25% BASE	N020330 001	Nov 04, 1993
AB	!	EQ 0.5% BASE	N020330 002	Nov 04, 1993

SOLUTION/DROPS; OPHTHALMIC

TIMOLOL MALEATE

AT	BAUSCH AND LOMB	EQ 0.25% BASE	A074778 001	Mar 25, 1997
AT	FDC LTD	EQ 0.25% BASE	A077259 001	Apr 30, 2008
AT	PACIFIC PHARMA	EQ 0.25% BASE	A074746 001	Mar 25, 1997
AT	! SANDOZ INC	EQ 0.25% BASE	A074261 001	Apr 28, 1995
AT	WOCKHARDT	EQ 0.25% BASE	A078771 001	Sep 28, 2009

TIMOPTIC

AT	+ VALEANT PHARMS INTL	EQ 0.25% BASE	N018086 001	
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TIMOLOL MALEATE

AT1	AKORN	EQ 0.5% BASE	A074466 001	Mar 25, 1997
AT1		EQ 0.5% BASE	A074516 001	Mar 25, 1997
AT1	BAUSCH AND LOMB	EQ 0.5% BASE	A074776 001	Mar 25, 1997
AT1	FDC LTD	EQ 0.5% BASE	A077259 002	Apr 30, 2008
AT1	HI TECH PHARMA	EQ 0.5% BASE	A075163 001	Sep 10, 2002
AT1	PACIFIC PHARMA	EQ 0.5% BASE	A074747 001	Mar 25, 1997
AT1	! SANDOZ INC	EQ 0.5% BASE	A074262 001	Apr 28, 1995
AT1	WOCKHARDT	EQ 0.5% BASE	A078771 002	Sep 28, 2009

TIMOPTIC

AT1	+ VALEANT PHARMS INTL	EQ 0.5% BASE	N018086 002	
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ISTALOL

AT2	! BAUSCH AND LOMB	EQ 0.5% BASE	N021516 001	Jun 04, 2004
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TIMOLOL MALEATE

AT2	APOTEX	EQ 0.5% BASE	A204936 001	Apr 17, 2015
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TIMOPTIC IN OCUDOSE

	! VALEANT PHARMS INTL	EQ 0.25% BASE	N019463 001	Nov 05, 1986
	!	EQ 0.5% BASE	N019463 002	Nov 05, 1986

TABLET; ORAL

TIMOLOL MALEATE

AB	ELYSIUM	5MG	A207556 001	May 02, 2019
AB		10MG	A207556 002	May 02, 2019
AB		20MG	A207556 003	May 02, 2019
AB	MYLAN	5MG	A072668 002	Jun 08, 1990
AB		10MG	A072668 003	Jun 08, 1990
AB	!	20MG	A072668 001	Jun 08, 1990

PRESCRIPTION DRUG PRODUCT LISTTINIDAZOLE

TABLET;ORAL

TINDAMAX

AB	+	MISSION PHARMA	250MG	N021618 001	May 17, 2004
AB	+	!	500MG	N021618 002	May 17, 2004

TINIDAZOLE

AB		EDENBRIDGE PHARMS	250MG	A203808 001	Aug 04, 2015
AB			500MG	A203808 002	Aug 04, 2015
AB		HIKMA	250MG	A201172 001	Apr 30, 2012
AB			500MG	A201172 002	Apr 30, 2012
AB		NOVEL LABS INC	250MG	A202044 001	Apr 30, 2012
AB			500MG	A202044 002	Apr 30, 2012
AB		UNIQUE PHARM LABS	250MG	A202489 001	Oct 09, 2013
AB			500MG	A202489 002	Oct 09, 2013

TIOPRONIN

TABLET;ORAL

THIOLA

+	!	MISSION PHARMA	100MG	N019569 001	Aug 11, 1988
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TABLET, DELAYED RELEASE;ORAL

THIOLA EC

+		MISSION PHARMACAL	100MG	N211843 001	Jun 28, 2019
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CO

+	!		300MG	N211843 002	Jun 28, 2019
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TIOTROPIUM BROMIDE

POWDER; INHALATION

SPIRIVA

+	!	BOEHRINGER INGELHEIM	EQ 0.018MG BASE/INH	N021395 001	Jan 30, 2004
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SPRAY, METERED; INHALATION

SPIRIVA RESPIMAT

+		BOEHRINGER INGELHEIM	EQ 0.00125MG BASE/INH	N021936 002	Sep 15, 2015
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+	!		EQ 0.0025MG BASE/INH	N021936 001	Sep 24, 2014
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TIPIRACIL HYDROCHLORIDE; TRIFLURIDINE

TABLET;ORAL

LONSURF

+		TAIHO ONCOLOGY	EQ 6.14MG BASE;15MG	N207981 001	Sep 22, 2015
+	!		EQ 8.19MG BASE;20MG	N207981 002	Sep 22, 2015

TIPRANAVIR

CAPSULE;ORAL

APTIVUS

+	!	BOEHRINGER INGELHEIM	250MG	N021814 001	Jun 22, 2005
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SOLUTION;ORAL

APTIVUS

+	!	BOEHRINGER INGELHEIM	100MG/ML	N022292 001	Jun 23, 2008
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TIROFIBAN HYDROCHLORIDE

INJECTABLE; INJECTION

AGGRASTAT

+		MEDICURE	EQ 5MG BASE/100ML (EQ 0.05MG BASE/ML)	N020913 002	May 17, 2002
+	!		EQ 12.5MG BASE/250ML (EQ 0.05MG BASE/ML)	N020913 003	Apr 20, 2000

SOLUTION; INJECTION

AGGRASTAT

+	!	MEDICURE	EQ 3.75MG BASE/15ML (EQ 0.25MG BASE/ML)	N020912 002	Aug 31, 2016
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TIZANIDINE HYDROCHLORIDE

CAPSULE;ORAL

TIZANIDINE HYDROCHLORIDE

AB		ALEMIC PHARMS LTD	EQ 2MG BASE	A213223 001	Jan 13, 2020
AB			EQ 4MG BASE	A213223 002	Jan 13, 2020
AB			EQ 6MG BASE	A213223 003	Jan 13, 2020
AB		ALKEM LABS LTD	EQ 2MG BASE	A212196 001	Mar 27, 2019
AB			EQ 4MG BASE	A212196 002	Mar 27, 2019
AB			EQ 6MG BASE	A212196 003	Mar 27, 2019
AB		APOTEX INC	EQ 2MG BASE	A078868 001	Feb 03, 2012
AB			EQ 4MG BASE	A078868 002	Feb 03, 2012
AB			EQ 6MG BASE	A078868 003	Feb 03, 2012
AB		CADILA PHARMS LTD	EQ 2MG BASE	A210021 001	Mar 26, 2019
AB			EQ 4MG BASE	A210021 002	Mar 26, 2019
AB			EQ 6MG BASE	A210021 003	Mar 26, 2019

PRESCRIPTION DRUG PRODUCT LIST

TIZANIDINE HYDROCHLORIDE

CAPSULE; ORAL

TIZANIDINE HYDROCHLORIDE

AB	JUBILANT GENERICS	<u>EQ 2MG BASE</u>	<u>A209605 001</u>	Aug 04, 2017
AB		<u>EQ 4MG BASE</u>	<u>A209605 002</u>	Aug 04, 2017
AB		<u>EQ 6MG BASE</u>	<u>A209605 003</u>	Aug 04, 2017
AB	MYLAN PHARMS INC	<u>EQ 2MG BASE</u>	<u>A091502 001</u>	Nov 09, 2012
AB		<u>EQ 4MG BASE</u>	<u>A091502 002</u>	Nov 09, 2012
AB		<u>EQ 6MG BASE</u>	<u>A091502 003</u>	Nov 09, 2012
AB	NOVAST LABS	<u>EQ 2MG BASE</u>	<u>A210267 001</u>	Mar 12, 2019
AB		<u>EQ 4MG BASE</u>	<u>A210267 002</u>	Mar 12, 2019
AB		<u>EQ 6MG BASE</u>	<u>A210267 003</u>	Mar 12, 2019
AB	PAR PHARM INC	<u>EQ 2MG BASE</u>	<u>A207199 001</u>	Mar 14, 2017
AB		<u>EQ 4MG BASE</u>	<u>A207199 002</u>	Mar 14, 2017
AB		<u>EQ 6MG BASE</u>	<u>A207199 003</u>	Mar 14, 2017
AB	ZYDUS PHARMS	<u>EQ 2MG BASE</u>	<u>A208622 001</u>	Mar 03, 2017
AB		<u>EQ 4MG BASE</u>	<u>A208622 002</u>	Mar 03, 2017
AB		<u>EQ 6MG BASE</u>	<u>A208622 003</u>	Mar 03, 2017

ZANAFLEX

AB	+ COVIS PHARMA BV	<u>EQ 2MG BASE</u>	<u>N021447 001</u>	Aug 29, 2002
AB	+	<u>EQ 4MG BASE</u>	<u>N021447 002</u>	Aug 29, 2002
AB	+!	<u>EQ 6MG BASE</u>	<u>N021447 003</u>	Aug 29, 2002

TABLET; ORAL

TIZANIDINE HYDROCHLORIDE

AB	ALKEM LABS LTD	<u>EQ 2MG BASE</u>	<u>A211798 001</u>	Jan 25, 2019
AB		<u>EQ 4MG BASE</u>	<u>A211798 002</u>	Jan 25, 2019
AB	APOTEX	<u>EQ 2MG BASE</u>	<u>A076533 001</u>	Jan 16, 2004
AB		<u>EQ 4MG BASE</u>	<u>A076533 002</u>	Jan 16, 2004
AB	CASI PHARMS INC	<u>EQ 2MG BASE</u>	<u>A076280 001</u>	Nov 26, 2002
AB		<u>EQ 4MG BASE</u>	<u>A076280 002</u>	Jun 27, 2002
AB	DR REDDYS LABS INC	<u>EQ 2MG BASE</u>	<u>A076286 001</u>	Jul 03, 2002
AB		<u>EQ 4MG BASE</u>	<u>A076286 002</u>	Jul 03, 2002
AB	EPIC PHARMA LLC	<u>EQ 2MG BASE</u>	<u>A076347 001</u>	Oct 11, 2002
AB		<u>EQ 4MG BASE</u>	<u>A076347 002</u>	Oct 11, 2002
AB	MYLAN	<u>EQ 2MG BASE</u>	<u>A076354 001</u>	Mar 28, 2003
AB		<u>EQ 4MG BASE</u>	<u>A076354 002</u>	Mar 28, 2003
AB	OXFORD PHARMS	<u>EQ 2MG BASE</u>	<u>A076281 001</u>	Oct 20, 2003
AB		<u>EQ 4MG BASE</u>	<u>A076281 002</u>	Oct 20, 2003
AB	SUN PHARM INDS INC	<u>EQ 2MG BASE</u>	<u>A076416 001</u>	Sep 29, 2003
AB		<u>EQ 4MG BASE</u>	<u>A076416 002</u>	Sep 29, 2003
AB	UNICHEM LABS LTD	<u>EQ 2MG BASE</u>	<u>A091283 001</u>	Nov 28, 2012
AB		<u>EQ 4MG BASE</u>	<u>A091283 002</u>	Nov 28, 2012
AB	ZYDUS PHARMS	<u>EQ 2MG BASE</u>	<u>A208187 001</u>	Mar 09, 2018
AB		<u>EQ 4MG BASE</u>	<u>A208187 002</u>	Mar 09, 2018

ZANAFLEX

AB	+! COVIS PHARMA BV	<u>EQ 4MG BASE</u>	<u>N020397 001</u>	Nov 27, 1996
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TOBRAMYCIN

OINTMENT; OPHTHALMIC

TOBEX

+! NOVARTIS

0.3%

N050555 001

POWDER; INHALATION

TOBI PODHALER

+! MYLAN SPECIALITY LP

28MG

N201688 001 Mar 22, 2013

SOLUTION; INHALATION

KITABIS PAK

AN	PULMOFLOW INC	<u>300MG/5ML</u>	<u>N205433 001</u>	Dec 02, 2014
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TOBI

AN	+! MYLAN SPECIALITY LP	<u>300MG/5ML</u>	<u>N050753 001</u>	Dec 22, 1997
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TOBRAMYCIN

AN	AKORN INC	<u>300MG/5ML</u>	<u>A201422 001</u>	May 28, 2014
AN	AMNEAL PHARMS	<u>300MG/5ML</u>	<u>A205501 001</u>	Jul 13, 2015
AN	DR REDDYS LABS SA	<u>300MG/5ML</u>	<u>A207080 001</u>	Jul 09, 2018
AN	LUPIN	<u>300MG/5ML</u>	<u>A208964 001</u>	Mar 22, 2017
AN	SUN PHARM	<u>300MG/5ML</u>	<u>A207136 001</u>	Dec 26, 2019
AN	TEVA PHARMS USA	<u>300MG/5ML</u>	<u>A091589 001</u>	Oct 10, 2013

BETHKIS

+! CHIESI USA INC

300MG/4ML

N201820 001 Oct 12, 2012

SOLUTION/DROPS; OPHTHALMIC

AKTOB

AT	AKORN	<u>0.3%</u>	<u>A064096 001</u>	Jan 31, 1996
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PRESCRIPTION DRUG PRODUCT LIST

TOBRAMYCIN

SOLUTION/DROPS;OPHTHALMIC

TOBRAMYCIN

<u>AT</u>	ALEMBIC PHARMS LTD	<u>0.3%</u>	<u>A211847</u>	<u>001</u>	Apr 19, 2019
<u>AT</u>	BAUSCH AND LOMB	<u>0.3%</u>	<u>A064052</u>	<u>001</u>	Nov 29, 1993
<u>AT</u>	FERA PHARMS	<u>0.3%</u>	<u>A065026</u>	<u>001</u>	Sep 11, 2001
<u>AT</u>	SOMERSET THERAPS LLC	<u>0.3%</u>	<u>A207444</u>	<u>001</u>	Jun 28, 2017

TOBREX

<u>AT</u>	<u>+</u> ! NOVARTIS	<u>0.3%</u>	<u>N050541</u>	<u>001</u>	
<u>AT</u>	SANDOZ INC	<u>0.3%</u>	<u>A062535</u>	<u>001</u>	Dec 13, 1984

TOBRAMYCIN SULFATE

INJECTABLE; INJECTION

TOBRAMYCIN SULFATE

<u>AP</u>	AKORN	<u>EQ 40MG BASE/ML</u>	<u>A205179</u>	<u>001</u>	Sep 16, 2014
<u>AP</u>	BAXTER HLTHCARE CORP	<u>EQ 40MG BASE/ML</u>	<u>A206965</u>	<u>001</u>	Jul 01, 2016
<u>AP</u>	FRESENIUS KABI USA	<u>EQ 10MG BASE/ML</u>	<u>A065122</u>	<u>001</u>	Nov 29, 2002
<u>AP</u>	!	<u>EQ 40MG BASE/ML</u>	<u>A065122</u>	<u>002</u>	Nov 29, 2002
<u>AP</u>		<u>EQ 1.2GM BASE/VIAL</u>	<u>N050789</u>	<u>001</u>	Jul 13, 2004
<u>AP</u>	! HOSPIRA	<u>EQ 10MG BASE/ML</u>	<u>A063112</u>	<u>001</u>	Apr 30, 1991
<u>AP</u>		<u>EQ 40MG BASE/ML</u>	<u>A063111</u>	<u>001</u>	Apr 30, 1991
<u>AP</u>	MYLAN LABS LTD	<u>EQ 40MG BASE/ML</u>	<u>A065407</u>	<u>001</u>	Mar 11, 2008
<u>AP</u>	TEVA PHARMS USA	<u>EQ 40MG BASE/ML</u>	<u>A063100</u>	<u>001</u>	Jan 30, 1992
<u>AP</u>	WEST-WARD PHARMS INT	<u>EQ 40MG BASE/ML</u>	<u>A063117</u>	<u>001</u>	Apr 26, 1991
<u>AP</u>	XELLIA PHARMS APS	<u>EQ 1.2GM BASE/VIAL</u>	<u>A205685</u>	<u>001</u>	Sep 16, 2014
<u>AP</u>	! XGEN PHARMS	<u>EQ 1.2GM BASE/VIAL</u>	<u>A065013</u>	<u>001</u>	Aug 17, 2001
	TOBRAMYCIN SULFATE (PHARMACY BULK)				
	! FRESENIUS KABI USA	EQ 40MG BASE/ML	A065120	001	Nov 29, 2002
	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

TOFACITINIB CITRATE

TABLET; ORAL

XELJANZ

+	PF PRISM CV	EQ 5MG BASE	N203214	001	Nov 06, 2012
<u>+</u> !		EQ 10MG BASE	N203214	002	May 30, 2018

TABLET, EXTENDED RELEASE; ORAL

XELJANZ XR

<u>+</u> !	PFIZER INC	EQ 11MG BASE	N208246	001	Feb 23, 2016
+		EQ 22MG BASE	N208246	002	Dec 12, 2019

TOLAZAMIDE

TABLET; ORAL

TOLAZAMIDE

	MYLAN PHARMS INC	250MG	A070259	001	Jan 02, 1986
!		500MG	A070259	003	Mar 17, 1986

TOLBUTAMIDE

TABLET; ORAL

TOLBUTAMIDE

!	MYLAN PHARMS INC	500MG	A086445	001	
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TOLCAPONE

TABLET; ORAL

TASMAR

<u>AB</u>	<u>+</u> ! BAUSCH	<u>100MG</u>	<u>N020697</u>	<u>001</u>	Jan 29, 1998
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TOLCAPONE

<u>AB</u>	MAYNE PHARMA	<u>100MG</u>	<u>A210095</u>	<u>001</u>	Aug 01, 2019
<u>AB</u>	NOVAST LABS	<u>100MG</u>	<u>A208937</u>	<u>001</u>	Aug 07, 2018
<u>AB</u>	PAR PHARM INC	<u>100MG</u>	<u>A204584</u>	<u>001</u>	Mar 26, 2015

TOLMETIN SODIUM

CAPSULE; ORAL

TOLMETIN SODIUM

<u>AB</u>	MYLAN	<u>EQ 400MG BASE</u>	<u>A073393</u>	<u>001</u>	May 27, 1993
<u>AB</u>	! TEVA	<u>EQ 400MG BASE</u>	<u>A073290</u>	<u>001</u>	Nov 27, 1991

TABLET; ORAL

TOLMETIN SODIUM

!	MYLAN	EQ 600MG BASE	A074473	001	Aug 30, 1994
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PRESCRIPTION DRUG PRODUCT LIST

TOLTERODINE TARTRATE

CAPSULE, EXTENDED RELEASE;ORAL

DETROL LA

AB	+	PHARMACIA AND UPJOHN	2MG	N021228 001	Dec 22, 2000
AB	+	!	4MG	N021228 002	Dec 22, 2000

TOLTERODINE TARTRATE

AB		HETERO LABS LTD III	2MG	A206419 001	Dec 12, 2017
AB			4MG	A206419 002	Dec 12, 2017
AB		INVENTIA HLTHCARE	2MG	A204562 001	Aug 19, 2019
AB			4MG	A204562 002	Aug 19, 2019
AB		MYLAN	2MG	A201486 001	Oct 31, 2013
AB			4MG	A201486 002	Oct 31, 2013
AB		TEVA PHARMS USA	2MG	A079141 001	Nov 22, 2016
AB			4MG	A079141 002	Nov 22, 2016
AB		TORRENT	2MG	A203016 001	Aug 11, 2015
AB			4MG	A203016 002	Aug 11, 2015

TABLET;ORAL

DETROL

AB	+	PHARMACIA AND UPJOHN	1MG	N020771 001	Mar 25, 1998
AB	+	!	2MG	N020771 002	Mar 25, 1998

TOLTERODINE TARTRATE

AB		INVATECH	1MG	A210775 001	Dec 30, 2019
AB			2MG	A210775 002	Dec 30, 2019
AB		IVAX SUB TEVA PHARMS	1MG	A077006 001	Feb 23, 2015
AB			2MG	A077006 002	Feb 23, 2015
AB		MACLEODS PHARMS LTD	1MG	A203409 001	Aug 31, 2015
AB			2MG	A203409 002	Aug 31, 2015
AB		UNIQUE PHARM LABS	1MG	A204721 001	Jan 24, 2020
AB			2MG	A204721 002	Jan 24, 2020

TOLVAPTAN

TABLET;ORAL

JYNARQUE

	+	OTSUKA PHARM CO LTD	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

SAMSCA

	+	OTSUKA AMERICA PHARM	15MG	N022275 001	May 19, 2009
	+	!	30MG	N022275 002	May 19, 2009

TOPIRAMATE

CAPSULE;ORAL

TOPAMAX

AB	+	JANSSEN PHARMS	15MG	N020844 001	Oct 26, 1998
AB	+	!	25MG	N020844 002	Oct 26, 1998

TOPIRAMATE

AB		TEVA	15MG	A076575 001	Apr 17, 2009
AB			25MG	A076575 002	Apr 17, 2009
AB		WATSON LABS	15MG	A077868 001	Apr 15, 2009
AB			25MG	A077868 002	Apr 15, 2009
AB		ZYDUS PHARMS USA INC	15MG	A078877 001	Oct 14, 2009
AB			25MG	A078877 002	Oct 14, 2009

CAPSULE, EXTENDED RELEASE;ORAL

TOPIRAMATE

AB		ZYDUS PHARMS	25MG	A207382 001	Nov 24, 2017
AB			50MG	A207382 002	Nov 24, 2017
AB			100MG	A207382 003	Nov 24, 2017

TROKENDI XR

AB	+	SUPERNU PHARMS	25MG	N201635 001	Aug 16, 2013
AB	+		50MG	N201635 002	Aug 16, 2013
AB	+		100MG	N201635 003	Aug 16, 2013

QUDEXY XR

BC	+	UPSHER SMITH LABS	25MG	N205122 001	Mar 11, 2014
BC	+		50MG	N205122 002	Mar 11, 2014
BC	+		100MG	N205122 003	Mar 11, 2014
BC	+	!	200MG	N205122 005	Mar 11, 2014

PRESCRIPTION DRUG PRODUCT LIST

3-428 (of 453)

TOPIRAMATE

CAPSULE, EXTENDED RELEASE;ORAL

TROKENDI XR

BC +! SUPERNUS PHARMS 200MG N201635 004 Aug 16, 2013

QUDEXY XR

+ UPSHER SMITH LABS 150MG N205122 004 Mar 11, 2014

TABLET;ORAL

TOPAMAXAB + JANSSEN PHARMS 25MG N020505 004 Dec 24, 1996AB + 50MG N020505 005 Dec 24, 1996AB +! 100MG N020505 001 Dec 24, 1996AB + 200MG N020505 002 Dec 24, 1996TOPIRAMATEAB ACCORD HLTHCARE 25MG A076311 001 Mar 27, 2009AB 50MG A076311 002 Mar 27, 2009AB 100MG A076311 003 Mar 27, 2009AB 200MG A076311 004 Mar 27, 2009AB APOTEX INC 25MG A077733 001 Mar 27, 2009AB 50MG A077733 002 Mar 27, 2009AB 100MG A077733 003 Mar 27, 2009AB 200MG A077733 004 Mar 27, 2009AB AUROBINDO PHARMA 25MG A078462 001 Mar 27, 2009AB 50MG A078462 002 Mar 27, 2009AB 100MG A078462 003 Mar 27, 2009AB 200MG A078462 004 Mar 27, 2009AB CIPLA LTD 25MG A076343 001 Mar 27, 2009AB 50MG A076343 002 Mar 27, 2009AB 100MG A076343 003 Mar 27, 2009AB 200MG A076343 004 Mar 27, 2009AB GLENMARK GENERICS 25MG A077627 001 Mar 27, 2009AB 50MG A077627 002 Mar 27, 2009AB 100MG A077627 003 Mar 27, 2009AB 200MG A077627 004 Mar 27, 2009AB INVAGEN PHARMS 25MG A079162 001 Mar 27, 2009AB 50MG A079162 002 Mar 27, 2009AB 100MG A079162 003 Mar 27, 2009AB 200MG A079162 004 Mar 27, 2009AB SUN PHARM 25MG A090278 001 Mar 27, 2009AB 50MG A090278 002 Mar 27, 2009AB 100MG A090278 003 Mar 27, 2009AB 200MG A090278 004 Mar 27, 2009AB SUN PHARM INDS LTD 25MG A076327 001 Mar 27, 2009AB 100MG A076327 002 Mar 27, 2009AB 200MG A076327 003 Mar 27, 2009AB TEVA 25MG A076317 001 Mar 27, 2009AB 50MG A076317 002 Mar 27, 2009AB 100MG A076317 003 Mar 27, 2009AB 200MG A076317 004 Mar 27, 2009AB TORRENT PHARMS 25MG A079153 001 Mar 27, 2009AB 50MG A079153 002 Mar 27, 2009AB 100MG A079153 003 Mar 27, 2009AB 200MG A079153 004 Mar 27, 2009AB UNICHEM LABS LTD 25MG A090162 001 Mar 27, 2009AB 50MG A090162 002 Mar 27, 2009AB 100MG A090162 003 Mar 27, 2009AB 200MG A090162 004 Feb 19, 2013AB UPSHER SMITH LABS 25MG A078499 001 Jan 07, 2010AB 50MG A078499 002 Jan 07, 2010AB 100MG A078499 003 Jan 07, 2010AB 200MG A078499 004 Jan 07, 2010AB ZYDUS PHARMS USA 25MG A078235 001 Mar 27, 2009AB INC 50MG A078235 002 Mar 27, 2009AB 100MG A078235 003 Mar 27, 2009AB 200MG A078235 004 Mar 27, 2009TOPOTECAN HYDROCHLORIDE

CAPSULE;ORAL

HYCAMTIN

+ NOVARTIS EQ 0.25MG BASE N020981 001 Oct 11, 2007

+! EQ 1MG BASE N020981 002 Oct 11, 2007

PRESCRIPTION DRUG PRODUCT LIST

TOPOTECAN HYDROCHLORIDE

INJECTABLE; INJECTION

HYCAMTIN

<u>AP</u>	<u>+!</u>	<u>NOVARTIS</u>	<u>EQ 4MG BASE/VIAL</u>	<u>N020671 001</u>	May 28, 1996
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TOPOTECAN HYDROCHLORIDE

<u>AP</u>		<u>ACCORD HLTHCARE</u>	<u>EQ 4MG BASE/VIAL</u>	<u>A202351 001</u>	Jun 26, 2013
<u>AP</u>		<u>ACTAVIS TOTOWA</u>	<u>EQ 4MG BASE/VIAL</u>	<u>A090620 001</u>	Dec 02, 2010
<u>AP</u>		<u>CIPLA</u>	<u>EQ 4MG BASE/VIAL</u>	<u>A091199 001</u>	Dec 01, 2010
<u>AP</u>		<u>DR REDDYS LABS LTD</u>	<u>EQ 4MG BASE/VIAL</u>	<u>A201191 001</u>	Mar 09, 2011
<u>AP</u>		<u>FRESENIUS KABI USA</u>	<u>EQ 4MG BASE/VIAL</u>	<u>A091089 001</u>	Nov 29, 2010
<u>AP</u>		<u>MEITHEAL</u>	<u>EQ 4MG BASE/VIAL</u>	<u>A201166 001</u>	Aug 08, 2012
<u>AP</u>		<u>NOVAST LABS</u>	<u>EQ 4MG BASE/VIAL</u>	<u>A206962 001</u>	Nov 30, 2016
<u>AP</u>		<u>SAGENT PHARMS</u>	<u>EQ 4MG BASE/VIAL</u>	<u>A091284 001</u>	Jan 26, 2011

SOLUTION; INTRAVENOUS

TOPOTECAN HYDROCHLORIDE

<u>AP</u>		<u>ACCORD HLTHCARE</u>	<u>EQ 4MG BASE/4ML (EQ 1MG BASE/ML)</u>	<u>A204406 002</u>	Jul 06, 2017
<u>AP</u>	<u>+!</u>	<u>HOSPIRA INC</u>	<u>EQ 4MG BASE/4ML (EQ 1MG BASE/ML)</u>	<u>N200582 001</u>	Feb 02, 2011
<u>AP</u>		<u>MYLAN LABS LTD</u>	<u>EQ 4MG BASE/4ML (EQ 1MG BASE/ML)</u>	<u>A206074 001</u>	Nov 24, 2017
<u>AP</u>		<u>TEVA PHARMS USA</u>	<u>EQ 4MG BASE/4ML (EQ 1MG BASE/ML)</u>	<u>N022453 001</u>	Dec 20, 2012
		<u>ACCORD HLTHCARE</u>	<u>EQ 1MG BASE/ML (EQ 1MG BASE/ML)</u>	<u>A204406 001</u>	Jul 06, 2017

TOREMIFENE CITRATE

TABLET; ORAL

FARESTON

<u>AB</u>	<u>+!</u>	<u>KYOWA KIRIN</u>	<u>EQ 60MG BASE</u>	<u>N020497 001</u>	May 29, 1997
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TOREMIFENE CITRATE

<u>AB</u>		<u>RISING</u>	<u>EQ 60MG BASE</u>	<u>A208813 001</u>	Dec 04, 2018
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TORSEMIDE

TABLET; ORAL

DEMADEX

<u>AB</u>	<u>+</u>	<u>MYLAN SPECIALITY LP</u>	<u>5MG</u>	<u>N020136 001</u>	Aug 23, 1993
<u>AB</u>	<u>+</u>		<u>10MG</u>	<u>N020136 002</u>	Aug 23, 1993
<u>AB</u>	<u>+!</u>		<u>20MG</u>	<u>N020136 003</u>	Aug 23, 1993
<u>AB</u>	<u>+</u>		<u>100MG</u>	<u>N020136 004</u>	Aug 23, 1993

TORSEMIDE

<u>AB</u>		<u>AUROBINDO PHARMA</u>	<u>5MG</u>	<u>A078249 001</u>	Oct 17, 2007
<u>AB</u>			<u>10MG</u>	<u>A078249 002</u>	Oct 17, 2007
<u>AB</u>			<u>20MG</u>	<u>A078249 003</u>	Oct 17, 2007
<u>AB</u>			<u>100MG</u>	<u>A078249 004</u>	Oct 17, 2007
<u>AB</u>		<u>COREPHARMA</u>	<u>5MG</u>	<u>A076894 001</u>	May 31, 2005
<u>AB</u>			<u>10MG</u>	<u>A076894 002</u>	May 31, 2005
<u>AB</u>			<u>20MG</u>	<u>A076894 003</u>	May 31, 2005
<u>AB</u>			<u>100MG</u>	<u>A076894 004</u>	May 31, 2005
<u>AB</u>		<u>HETERO LABS LTD III</u>	<u>5MG</u>	<u>A079234 001</u>	Jan 27, 2009
<u>AB</u>			<u>10MG</u>	<u>A079234 002</u>	Jan 27, 2009
<u>AB</u>			<u>20MG</u>	<u>A079234 003</u>	Jan 27, 2009
<u>AB</u>			<u>100MG</u>	<u>A079234 004</u>	Jan 27, 2009
<u>AB</u>		<u>HIKMA</u>	<u>5MG</u>	<u>A076943 001</u>	Mar 01, 2005
<u>AB</u>			<u>10MG</u>	<u>A076943 002</u>	Mar 01, 2005
<u>AB</u>			<u>20MG</u>	<u>A076943 003</u>	Mar 01, 2005
<u>AB</u>		<u>PAR PHARM</u>	<u>5MG</u>	<u>A076226 001</u>	May 27, 2003
<u>AB</u>			<u>10MG</u>	<u>A076226 002</u>	May 27, 2003
<u>AB</u>			<u>20MG</u>	<u>A076226 003</u>	May 27, 2003
<u>AB</u>			<u>100MG</u>	<u>A076226 004</u>	May 27, 2003
<u>AB</u>		<u>PLIVA PHARM IND</u>	<u>5MG</u>	<u>A076346 001</u>	May 30, 2003
<u>AB</u>			<u>10MG</u>	<u>A076346 002</u>	May 30, 2003
<u>AB</u>			<u>20MG</u>	<u>A076346 003</u>	May 30, 2003
<u>AB</u>			<u>100MG</u>	<u>A076346 004</u>	Oct 19, 2004
<u>AB</u>		<u>TEVA</u>	<u>5MG</u>	<u>A076110 001</u>	May 14, 2002
<u>AB</u>			<u>10MG</u>	<u>A076110 002</u>	May 14, 2002
<u>AB</u>			<u>20MG</u>	<u>A076110 003</u>	May 14, 2002
<u>AB</u>			<u>100MG</u>	<u>A076110 004</u>	May 14, 2002
<u>AB</u>		<u>VINTAGE PHARMS</u>	<u>5MG</u>	<u>A090613 001</u>	Mar 22, 2011
<u>AB</u>			<u>10MG</u>	<u>A090613 002</u>	Mar 22, 2011
<u>AB</u>			<u>20MG</u>	<u>A090613 003</u>	Mar 22, 2011
<u>AB</u>			<u>100MG</u>	<u>A090613 004</u>	Mar 22, 2011

PRESCRIPTION DRUG PRODUCT LIST

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
+		150MG	N022370	004	Aug 01, 2011
+		200MG	N022370	002	May 07, 2010
+		300MG	N022370	003	May 07, 2010

TABLET; ORAL

TRAMADOL HYDROCHLORIDE

AB	ACI HEALTHCARE LTD	50MG	A202075	001	Nov 28, 2011
AB	AMNEAL PHARMS	50MG	A076003	001	Jun 20, 2002
AB	APOTEX	50MG	A075981	001	Jul 10, 2002
AB	AUROBINDO PHARMA LTD	50MG	A203494	001	Mar 31, 2014
AB	CSPEC OUYI PHARM CO	50MG	A091498	001	Mar 29, 2013
AB	IPCA LABS LTD	50MG	A201973	001	Nov 16, 2012
AB	MACLEODS PHARMS LTD	50MG	A205702	001	Sep 25, 2015
AB	MYLAN	50MG	A075986	001	Jun 21, 2002
AB	PLIVA	50MG	A075982	001	Jul 01, 2002
AB	RUBICON	50MG	A208708	001	Jun 28, 2019
AB		100MG	A208708	002	Jun 28, 2019
AB	SUN PHARM INDS INC	50MG	A075964	001	Jun 19, 2002
AB	TEVA	50MG	A075977	001	Jun 19, 2002
AB	UNICHEM LABS LTD	50MG	A211825	001	Aug 09, 2019
AB	ZYDUS PHARMS USA INC	50MG	A090404	001	Jan 31, 2011

ULTRAM

AB	+!	JANSSEN PHARMS	50MG	N020281	002	Mar 03, 1995
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TABLET, EXTENDED RELEASE; ORAL

TRAMADOL HYDROCHLORIDE

AB1	!	LUPIN LTD	100MG	A200503	001	Aug 29, 2011
AB1			200MG	A200503	002	Aug 29, 2011
AB1			300MG	A200503	003	Aug 29, 2011
AB1		MYLAN	100MG	A205257	001	Dec 22, 2015
AB1			200MG	A205257	002	Dec 22, 2015
AB1			300MG	A205257	003	Dec 22, 2015
AB1		SUN PHARM	100MG	A201384	001	Dec 07, 2011
AB1			200MG	A201384	002	Dec 07, 2011
AB1			300MG	A201384	003	Dec 07, 2011
AB2		ACTAVIS ELIZABETH	100MG	A091609	001	Jun 27, 2012
AB2			200MG	A091609	002	Jun 27, 2012
AB2			300MG	A091609	003	Jun 27, 2012
AB2		ANCHEN PHARMS	100MG	A200491	001	Jun 27, 2012
AB2			200MG	A200491	002	Jun 27, 2012
AB2			300MG	A200491	003	Jun 27, 2012
AB2	!	SUN PHARM	100MG	A091607	001	Dec 30, 2011
AB2			200MG	A091607	002	Dec 30, 2011
AB2			300MG	A091607	003	Dec 30, 2011

TRAMETINIB DIMETHYL SULFOXIDE

TABLET; ORAL

MEKINIST

+	NOVARTIS	EQ 0.5MG	N204114	001	May 29, 2013
+!		EQ 2MG	N204114	003	May 29, 2013

TRANDOLAPRIL

TABLET; ORAL

TRANDOLAPRIL

AB		AUROBINDO PHARMA	1MG	A078438	001	Jun 12, 2007
AB			2MG	A078438	002	Jun 12, 2007
AB			4MG	A078438	003	Jun 12, 2007
AB		EPIC PHARMA	1MG	A078508	003	Jun 18, 2008
AB			2MG	A078508	001	Jun 18, 2008
AB			4MG	A078508	002	Jun 18, 2008
AB		LUPIN	1MG	A077522	001	Jun 12, 2007
AB			2MG	A077522	002	Jun 12, 2007
AB	!		4MG	A077522	003	Jun 12, 2007
AB		TEVA PHARMS	1MG	A077489	001	Dec 12, 2006
AB			2MG	A077489	002	Dec 12, 2006

PRESCRIPTION DRUG PRODUCT LISTTRANDOLAPRIL

TABLET; ORAL

TRANDOLAPRIL**AB** **+** **4MG** **A077489 003** Dec 12, 2006TRANDOLAPRIL; VERAPAMIL HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL

TARKA**AB** **+** ABEVIE **2MG;180MG** **N020591 001** Oct 22, 1996**AB** **+** **2MG;240MG** **N020591 004** Oct 22, 1996**AB** **+**! **4MG;240MG** **N020591 002** Oct 22, 1996TRANDOLAPRIL AND VERAPAMIL HYDROCHLORIDE**AB** GLENMARK GENERICS **2MG;180MG** **A079135 001** May 26, 2010**AB** **2MG;240MG** **A079135 002** May 26, 2010**AB** **4MG;240MG** **A079135 003** May 05, 2010**1MG;240MG** **A079135 004** Aug 30, 2010TRANEXAMIC ACID

INJECTABLE; INJECTION

CYKLOKAPRON**AP** **+**! PHARMACIA AND UPJOHN **100MG/ML** **N019281 001** Dec 30, 1986TRANEXAMIC ACID**AP** ACIC PHARMS **100MG/ML** **A202436 001** Feb 11, 2014**AP** AKORN **100MG/ML** **A202373 001** Nov 17, 2011**AP** **100MG/ML** **A206594 001** Sep 28, 2017**AP** **100MG/ML** **A206634 001** Jun 09, 2016**AP** AM REGENT **100MG/ML** **A201885 001** Aug 10, 2011**AP** AMNEAL PHARMS CO **100MG/ML** **A208840 001** Feb 28, 2017**AP** APOLLO **100MG/ML** **A212676 001** Jul 17, 2019**AP** APOTEX **100MG/ML** **A209860 001** Jan 14, 2020**AP** AUROBINDO PHARMA LTD **100MG/ML** **A205035 001** Jan 14, 2016**AP** EMCURE PHARMS LTD **100MG/ML** **A203521 001** Aug 12, 2014**AP** FRESENIUS KABI USA **100MG/ML** **A091596 001** Mar 02, 2012**AP** GLAND PHARMA LTD **100MG/ML** **A207239 001** Feb 13, 2017**AP** MICRO LABS **100MG/ML** **A206713 001** Jun 27, 2017**AP** MYLAN INSTITUTIONAL **100MG/ML** **A091657 001** Nov 03, 2011**AP** XGEN PHARMS **100MG/ML** **A201580 001** Jun 14, 2013**AP** ZYDUS PHARMS **100MG/ML** **A205228 001** Jul 17, 2017

SOLUTION; INTRAVENOUS

TRANEXAMIC ACID**+**! EXELA PHARMA SCS LLC **1GM/100ML (10MG/ML)** **N212020 001** Apr 15, 2019

TABLET; ORAL

LYSTEDA**AB** **+**! FERRING PHARMS INC **650MG** **N022430 001** Nov 13, 2009TRANEXAMIC ACID**AB** ACTAVIS LABS FL INC **650MG** **A202093 001** Dec 27, 2012**AB** APOTEX INC **650MG** **A202286 001** Jan 27, 2014**AB** MYLAN **650MG** **A205133 001** Sep 21, 2015TRANLYCYPROMINE SULFATE

TABLET; ORAL

PARNATE**AB** **+**! CONCORDIA **EQ 10MG BASE** **N012342 003** Aug 16, 1985TRANLYCYPROMINE SULFATE**AB** CNTY LINE PHARMS **EQ 10MG BASE** **A206856 001** Apr 17, 2018**AB** PAR PHARM **EQ 10MG BASE** **A040640 001** Jun 29, 2006TRAVOPROST

SOLUTION/DROPS; OPHTHALMIC

TRAVATAN Z**AT** **+**! NOVARTIS **0.004%** **N021994 001** Sep 21, 2006TRAVOPROST**AT** APOTEX **0.004%** **A203431 001** Jul 10, 2015**AT** MYLAN **0.004%** **A205050 001** Jul 07, 2017**AT1** ALEMBIC PHARMS LTD **0.004%** **A210458 001** Dec 20, 2019**AT1** ! PAR PHARM **0.004%** **A091340 001** Mar 01, 2013

PRESCRIPTION DRUG PRODUCT LIST

TRAZODONE HYDROCHLORIDE

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	ALVOGEN	50MG	A071636 001	Apr 18, 1988
AB		100MG	A071514 001	Apr 18, 1988
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	AUROLIFE PHARMA LLC	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	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

TREPROSTINIL

INJECTABLE; IV (INFUSION), SUBCUTANEOUS

REMODYLIN

AP	! UNITED THERAP	1MG/ML	N021272 001	May 21, 2002
AP	!	2.5MG/ML	N021272 002	May 21, 2002
AP	!	5MG/ML	N021272 003	May 21, 2002
AP	!	10MG/ML	N021272 004	May 21, 2002

TREPROSTINIL

AP	PAR STERILE PRODUCTS	1MG/ML	A209382 001	Sep 24, 2019
AP		2.5MG/ML	A209382 002	Sep 24, 2019
AP		5MG/ML	A209382 003	Sep 24, 2019
AP		10MG/ML	A209382 004	Sep 24, 2019
AP	SANDOZ INC	1MG/ML	A203649 001	Nov 30, 2017
AP		2.5MG/ML	A203649 002	Nov 30, 2017
AP		5MG/ML	A203649 003	Nov 30, 2017
AP		10MG/ML	A203649 004	Nov 30, 2017
AP	TEVA PHARMS USA	1MG/ML	A206648 001	Sep 26, 2019
AP		2.5MG/ML	A206648 002	Sep 26, 2019
AP		5MG/ML	A206648 003	Sep 26, 2019
AP		10MG/ML	A206648 004	Sep 26, 2019

SOLUTION; INHALATION

TYVASO

! UNITED THERAP	0.6MG/ML	N022387 001	Jul 30, 2009
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SOLUTION; INTRAVENOUS, SUBCUTANEOUS

REMODYLIN

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

TREPROSTINIL DIOLAMINE

TABLET, EXTENDED RELEASE; ORAL

ORENITRAM

+	UNITED THERAP	EQ 0.125MG BASE	N203496 001	Dec 20, 2013
+		EQ 0.25MG BASE	N203496 002	Dec 20, 2013
!		EQ 1MG BASE	N203496 003	Dec 20, 2013
+		EQ 2.5MG BASE	N203496 004	Dec 20, 2013
+		EQ 5MG BASE	N203496 005	Oct 07, 2016

PRESCRIPTION DRUG PRODUCT LIST

3-433 (of 453)

TRETINOIN

CAPSULE;ORAL

TRETINOIN

AB	ANCHEN PHARMS	10MG	A201687 001	Oct 24, 2012
AB	! BARR LABS INC	10MG	A077684 001	Jun 22, 2007
AB	GLENMARK PHARMS LTD	10MG	A208279 001	Dec 23, 2016

CREAM;TOPICAL

AVITA

AB	MYLAN PHARMS INC	0.025%	N020404 003	Jan 14, 1997
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RETIN-A

AB	+! VALEANT BERMUDA	0.025%	N019049 001	Sep 16, 1988
AB	+! VALEANT PHARMS NORTH	0.1%	N017340 001	

TRETINOIN

AB	PERRIGO PHARMA INTL	0.025%	A075264 001	Dec 24, 1998
AB		0.1%	A075213 001	Dec 24, 1998
AB	TARO PHARMS	0.1%	A211645 001	Jan 22, 2019

RETIN-A

AB1	+! VALEANT BERMUDA	0.05%	N017522 001	
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TRETINOIN

AB1	PERRIGO PHARMA INTL	0.05%	A075265 001	Dec 24, 1998
AB1	TARO PHARMS	0.05%	A211644 001	Jan 25, 2019

RENOVA

AB2	+! VALEANT PHARMS NORTH	0.05%	N019963 001	Dec 29, 1995
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TRETINOIN

AB2	ZO SKIN HEALTH	0.05%	A076498 001	Sep 15, 2005
	RENOVA			
	+! VALEANT PHARMS NORTH	0.02%	N021108 001	Aug 31, 2000

GEL;TOPICAL

ATRALIN

AB	+! DOW PHARM	0.05%	N022070 001	Jul 26, 2007
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RETIN-A

AB	+! VALEANT INTL	0.01%	N017955 001	
AB	+!	0.025%	N017579 002	

RETIN-A MICRO

AB	+! VALEANT INTL	0.04%	N020475 002	May 10, 2002
AB	+!	0.1%	N020475 001	Feb 07, 1997

TRETINOIN

AB	MYLAN	0.04%	A202567 001	Jul 17, 2013
AB		0.05%	A207955 001	Aug 13, 2015
AB		0.1%	A202026 001	Jul 17, 2013
AB	PERRIGO PHARMA INTL	0.01%	A075589 001	Jun 11, 2002
AB		0.025%	A075529 001	Feb 22, 2000

AVITA

BT	MYLAN	0.025%	N020400 001	Jan 29, 1998
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RETIN-A-MICRO

	+! VALEANT INTL	0.06%	N020475 004	Oct 23, 2017
	+!	0.08%	N020475 003	Jan 28, 2014

LOTION;TOPICAL

ALTRENO

	+! DOW PHARM	0.05%	N209353 001	Aug 23, 2018
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TRIAMCINOLONE ACETONIDE

CREAM;TOPICAL

TRIAMCINOLONE ACETONIDE

AT	ACP NIMBLE	0.025%	A089797 001	May 31, 1991
AT		0.1%	A089798 001	May 31, 1991
AT	ALKEM LABS LTD	0.025%	A207651 001	Dec 26, 2017
AT		0.1%	A207651 002	Dec 26, 2017
AT		0.5%	A207651 003	Dec 26, 2017
AT	! FOUGERA PHARMS	0.025%	A085692 001	
AT	!	0.1%	A085692 003	
AT	!	0.5%	A085692 002	
AT	GLENMARK PHARMS LTD	0.1%	A207117 001	Aug 05, 2016
AT	LUPIN ATLANTIS	0.025%	A208763 001	Feb 01, 2017
AT		0.1%	A208763 002	Feb 01, 2017
AT		0.5%	A208763 003	Feb 01, 2017
AT	MACLEODS PHARMS LTD	0.025%	A209535 001	May 18, 2018
AT		0.1%	A209535 002	May 18, 2018
AT		0.5%	A209535 003	May 18, 2018
AT	MLV	0.025%	A040671 001	Jun 09, 2006

PRESCRIPTION DRUG PRODUCT LIST

TRIAMCINOLONE ACETONIDE

CREAM; TOPICAL

TRIAMCINOLONE ACETONIDE

<u>AT</u>		<u>0.1%</u>	<u>A040671 002</u>	Jun 09, 2006
<u>AT</u>	+	<u>0.025%</u>	<u>N011601 003</u>	
<u>AT</u>	+	<u>0.1%</u>	<u>N011601 006</u>	
<u>AT</u>		<u>0.025%</u>	<u>A086413 002</u>	
<u>AT</u>		<u>0.1%</u>	<u>A086413 003</u>	
<u>AT</u>		<u>0.5%</u>	<u>A086413 001</u>	
<u>AT</u>	STRIDES PHARMA	<u>0.025%</u>	<u>A210346 001</u>	Feb 11, 2019
<u>AT</u>		<u>0.1%</u>	<u>A210346 002</u>	Feb 11, 2019
<u>AT</u>		<u>0.5%</u>	<u>A210346 003</u>	Feb 11, 2019
<u>AT</u>	TARO PHARM INDS LTD	<u>0.1%</u>	<u>A040039 001</u>	Nov 26, 1997
<u>AT</u>	TELOGENT PHARMA INC	<u>0.1%</u>	<u>A208848 001</u>	Sep 18, 2017

TRIDERM

<u>AT</u>	CROWN LABS	<u>0.025%</u>	<u>A088042 002</u>	Mar 25, 2015
<u>AT</u>		<u>0.1%</u>	<u>A088042 001</u>	Mar 19, 1984
<u>AT</u>		<u>0.5%</u>	<u>A088042 003</u>	Mar 25, 2015

FOR SUSPENSION, EXTENDED RELEASE; INTRA-ARTICULAR
ZILRETTA

+	FLEXION THERAPS INC	32MG/VIAL	N208845 001	Oct 06, 2017
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INJECTABLE; INJECTION

KENALOG-40

<u>AB</u>	+	APOTHECON	<u>40MG/ML</u>	<u>N014901 001</u>	
<u>AB</u>		AMNEAL	<u>40MG/ML</u>	<u>A207550 001</u>	Dec 11, 2017
<u>AB</u>		TEVA PHARMS USA	<u>40MG/ML</u>	<u>A209852 001</u>	Oct 05, 2018
		KENALOG-10			
	+	APOTHECON	10MG/ML	N012041 001	
		KENALOG-80			
	+	APOTHECON	80MG/ML	N014901 002	Apr 12, 2019

INJECTABLE; INTRAVITREAL

TRIESENCE

+	NOVARTIS	40MG/ML (40MG/ML)	N022048 001	Nov 29, 2007
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LOTION; TOPICAL

TRIAMCINOLONE ACETONIDE

<u>AT</u>	ACP NIMBLE	<u>0.1%</u>	<u>A089129 001</u>	Aug 14, 1986	
<u>AT</u>	AKORN	<u>0.025%</u>	<u>A202374 001</u>	May 08, 2013	
<u>AT</u>		<u>0.1%</u>	<u>A202374 002</u>	May 08, 2013	
<u>AT</u>	ALLIED	<u>0.1%</u>	<u>A040672 002</u>	Dec 13, 2006	
<u>AT</u>	FOUGERA PHARMS	<u>0.025%</u>	<u>A040467 001</u>	Apr 21, 2003	
<u>AT</u>		<u>0.1%</u>	<u>A040467 002</u>	Apr 21, 2003	
<u>AT</u>	TELOGENT PHARMA INC	<u>0.025%</u>	<u>A204608 001</u>	Jul 07, 2016	
<u>AT</u>		<u>0.1%</u>	<u>A204606 001</u>	Jul 07, 2016	
<u>AT</u>	!	WOCKHARDT BIO AG	<u>0.025%</u>	<u>A088450 001</u>	Apr 01, 1985
<u>AT</u>	!		<u>0.1%</u>	<u>A088451 001</u>	Apr 03, 1985

OINTMENT; TOPICAL

TRIAMCINOLONE ACETATE

<u>AT</u>	MACLEODS PHARMS LTD	<u>0.025%</u>	<u>A209828 001</u>	Nov 23, 2018
<u>AT</u>		<u>0.1%</u>	<u>A209828 002</u>	Nov 23, 2018
<u>AT</u>		<u>0.5%</u>	<u>A209828 003</u>	Nov 23, 2018

TRIAMCINOLONE ACETONIDE

<u>AT</u>	ACP NIMBLE	<u>0.025%</u>	<u>A089795 001</u>	Dec 23, 1988
<u>AT</u>		<u>0.1%</u>	<u>A089796 001</u>	Dec 23, 1988
<u>AT</u>		<u>0.5%</u>	<u>A208925 001</u>	Oct 06, 2017
<u>AT</u>	ENCUBE	<u>0.05%</u>	<u>A212384 001</u>	Nov 29, 2019
<u>AT</u>	FOUGERA PHARMS	<u>0.025%</u>	<u>A085691 001</u>	
<u>AT</u>		<u>0.1%</u>	<u>A085691 003</u>	
<u>AT</u>		<u>0.5%</u>	<u>A085691 002</u>	
<u>AT</u>	GLENMARK PHARMS	<u>0.1%</u>	<u>A208320 001</u>	Aug 22, 2017
<u>AT</u>	GLENMARK PHARMS LTD	<u>0.5%</u>	<u>A206379 001</u>	Jul 22, 2016
<u>AT</u>	NOVEL LABS INC	<u>0.1%</u>	<u>A207365 001</u>	Oct 12, 2018
<u>AT</u>	!	PERRIGO NEW YORK	<u>0.025%</u>	<u>A087385 002</u>
<u>AT</u>	!		<u>0.1%</u>	<u>A087385 003</u>
<u>AT</u>	!		<u>0.5%</u>	<u>A087385 001</u>
<u>AT</u>	STRIDES PHARMA	<u>0.1%</u>	<u>A211315 001</u>	Mar 18, 2020
<u>AT</u>		<u>0.5%</u>	<u>A211315 002</u>	Mar 18, 2020
<u>AT</u>	TARO PHARM INDS LTD	<u>0.1%</u>	<u>A040037 001</u>	Sep 30, 1994
<u>AT</u>	TELOGENT PHARMA INC	<u>0.1%</u>	<u>A205373 001</u>	May 13, 2016
<u>AT</u>		<u>0.5%</u>	<u>A208590 001</u>	Mar 03, 2017

TRIAMCINOLONE ACETONIDE IN ABSORBASE

<u>AT</u>	!	CMP PHARMA INC	<u>0.05%</u>	<u>A089595 001</u>	Mar 23, 1995
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PRESCRIPTION DRUG PRODUCT LIST

TRIAMCINOLONE ACETONIDE

PASTE; DENTAL

TRIAMCINOLONE ACETONIDE

<u>AT</u>	ACP NIMBLE	<u>0.1%</u>	<u>A205592</u>	<u>001</u>	Jan 12, 2017
<u>AT</u>	AKORN	<u>0.1%</u>	<u>A206312</u>	<u>001</u>	Aug 11, 2016
<u>AT</u>	LYNE	<u>0.1%</u>	<u>A040771</u>	<u>001</u>	Jul 01, 2010
<u>AT</u>	! TARO	<u>0.1%</u>	<u>A070730</u>	<u>001</u>	Oct 01, 1986

SPRAY; TOPICAL

KENALOG

<u>AT</u>	+! SUN PHARM INDS INC	<u>0.147MG/GM</u>	<u>N012104</u>	<u>001</u>	
<u>TRIAMCINOLONE ACETONIDE</u>					
<u>AT</u>	AKORN	<u>0.147MG/GM</u>	<u>A207094</u>	<u>001</u>	Dec 07, 2016
<u>AT</u>	PERRIGO UK FINCO	<u>0.147MG/GM</u>	<u>A205782</u>	<u>001</u>	Apr 13, 2015
<u>AT</u>	RISING	<u>0.147MG/GM</u>	<u>A206786</u>	<u>001</u>	Sep 08, 2017

TRIAMCINOLONE HEXACETONIDE

INJECTABLE; INJECTION

ARISTOSPAN

	+! SANDOZ INC	5MG/ML	N016466	001	
	+!	20MG/ML	N016466	002	

TRIAMTERENE

CAPSULE; ORAL

DYRENIUM

<u>AB</u>	+ CONCORDIA	<u>50MG</u>	<u>N013174</u>	<u>001</u>	
<u>AB</u>	+!	<u>100MG</u>	<u>N013174</u>	<u>002</u>	
<u>TRIAMTERENE</u>					
<u>AB</u>	AGNITIO	<u>50MG</u>	<u>A211581</u>	<u>001</u>	Aug 19, 2019
<u>AB</u>		<u>100MG</u>	<u>A211581</u>	<u>002</u>	Aug 19, 2019

TRIAZOLAM

TABLET; ORAL

HALCION

<u>AB</u>	+ PHARMACIA AND UPJOHN	<u>0.125MG</u>	<u>N017892</u>	<u>003</u>	Apr 26, 1985
<u>AB</u>	+!	<u>0.25MG</u>	<u>N017892</u>	<u>001</u>	Nov 15, 1982
<u>TRIAZOLAM</u>					
<u>AB</u>	HIKMA	<u>0.125MG</u>	<u>A074224</u>	<u>001</u>	Jun 01, 1994
<u>AB</u>		<u>0.25MG</u>	<u>A074224</u>	<u>002</u>	Jun 01, 1994

TRICLABENDAZOLE

TABLET; ORAL

EGATEN

	+! NOVARTIS	250MG	N208711	001	Feb 13, 2019
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TRIENTINE HYDROCHLORIDE

CAPSULE; ORAL

CLOVIQUE

<u>AB</u>	KADMON PHARMS LLC	<u>250MG</u>	<u>A209731</u>	<u>001</u>	Oct 21, 2019
<u>SYPRINE</u>					
<u>AB</u>	+! BAUSCH	<u>250MG</u>	<u>N019194</u>	<u>001</u>	Nov 08, 1985

TRIENTINE HYDROCHLORIDE

<u>AB</u>	AMNEAL PHARMS LLC	<u>250MG</u>	<u>A210619</u>	<u>001</u>	Feb 08, 2019
<u>AB</u>	DR REDDYS LABS LTD	<u>250MG</u>	<u>A211076</u>	<u>001</u>	Jul 03, 2019
<u>AB</u>	KADMON PHARMS LLC	<u>250MG</u>	<u>A209415</u>	<u>001</u>	Sep 16, 2019
<u>AB</u>	MSN	<u>250MG</u>	<u>A211134</u>	<u>001</u>	May 22, 2019
<u>AB</u>	NAVINTA LLC	<u>250MG</u>	<u>A211251</u>	<u>001</u>	Jan 16, 2019
<u>AB</u>	PAR PHARM INC	<u>250MG</u>	<u>A210096</u>	<u>001</u>	Sep 25, 2019
<u>AB</u>	RISING	<u>250MG</u>	<u>A212238</u>	<u>001</u>	Feb 20, 2020
<u>AB</u>	WATSON LABS TEVA	<u>250MG</u>	<u>A207567</u>	<u>001</u>	Feb 07, 2018
<u>AB</u>	ZYDUS PHARMS	<u>250MG</u>	<u>A211554</u>	<u>001</u>	Apr 26, 2019

TRIFAROTENE

CREAM; TOPICAL

AKLIEF

	+! GALDERMA R AND D	0.005%	N211527	001	Oct 04, 2019
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TRIFLUOPERAZINE HYDROCHLORIDE

TABLET; ORAL

TRIFLUOPERAZINE HYDROCHLORIDE

<u>AB</u>	MYLAN	<u>EQ 1MG BASE</u>	<u>A040209</u>	<u>001</u>	Jul 07, 1997
<u>AB</u>		<u>EQ 2MG BASE</u>	<u>A040209</u>	<u>002</u>	Jul 07, 1997
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A040209</u>	<u>003</u>	Jul 07, 1997
<u>AB</u>	!	<u>EQ 10MG BASE</u>	<u>A040209</u>	<u>004</u>	Jul 07, 1997
<u>AB</u>	SANDOZ	<u>EQ 1MG BASE</u>	<u>A085785</u>	<u>001</u>	
<u>AB</u>		<u>EQ 2MG BASE</u>	<u>A085786</u>	<u>001</u>	

PRESCRIPTION DRUG PRODUCT LIST

TRIFLUOPERAZINE HYDROCHLORIDE

TABLET; ORAL

TRIFLUOPERAZINE HYDROCHLORIDE

AB		EQ 5MG BASE	A085789	001	
AB		EQ 10MG BASE	A085788	001	

TRIFLURIDINE

SOLUTION/DROPS; OPHTHALMIC

TRIFLURIDINE

AT	HI TECH	1%	A205438	001	Jul 28, 2017
AT	SANDOZ INC	1%	A074311	001	Oct 06, 1995
VIROPTIC					
AT	+!	MONARCH PHARMS	1%	N018299	001

TRIHEXYPHENIDYL HYDROCHLORIDE

ELIXIR; ORAL

TRIHEXYPHENIDYL HYDROCHLORIDE

AA	MIKART	2MG/5ML	A040251	001	Sep 27, 1999	
AA	!	PHARM ASSOC	2MG/5ML	A040177	001	Apr 17, 1997

TABLET; ORAL

TRIHEXYPHENIDYL 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

TRIMETHOBENZAMIDE HYDROCHLORIDE

CAPSULE; ORAL

TIGAN

AB	+!	KING PHARMS LLC	300MG	N017531	006	Dec 13, 2001
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TRIMETHOBENZAMIDE HYDROCHLORIDE

AB	LUPIN	300MG	A076546	001	Aug 20, 2003
AB	SUN PHARM INDUSTRIES	300MG	A076570	001	Aug 28, 2003

INJECTABLE; INJECTION

TIGAN

	+!	PAR STERILE PRODUCTS	100MG/ML	N017530	001	
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TRIMETHOPRIM

TABLET; ORAL

TRIMETHOPRIM

AB	+!	MAYNE PHARMA	100MG	N018679	001	Jul 30, 1982
AB		NOVEL LABS INC	100MG	A091437	001	Jun 15, 2011
AB		WATSON LABS	100MG	A070049	001	Jun 06, 1985

TRIMETHOPRIM HYDROCHLORIDE

SOLUTION; ORAL

PRIMSOL

	+!	ALLEGIS	EQ 50MG BASE/5ML	N074973	001	Jan 24, 2000
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TRIMIPRAMINE MALEATE

CAPSULE; ORAL

TRIMIPRAMINE MALEATE

AB		CROSSMEDIKA SA	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

	+!	ARBOR PHARMS LLC	EQ 22.5MG BASE/VIAL	N208956	001	Jun 29, 2017
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INJECTABLE; INTRAMUSCULAR

TRELSTAR

	+!	ALLERGAN	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

PRESCRIPTION DRUG PRODUCT LISTTROPICAMIDE

SOLUTION/DROPS;OPHTHALMIC

MYDRIACYL

AT	!	NOVARTIS	1%	A084306	001	
AT	!	SANDOZ INC	0.5%	A084305	001	

TROPICACYL

AT		AKORN	0.5%	A040314	001	Sep 29, 2000
AT			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

TROSPIUM CHLORIDE

CAPSULE, EXTENDED RELEASE;ORAL

TROSPIUM CHLORIDE

AB	!	ACTAVIS LABS FL INC	60MG	A091289	001	Oct 12, 2012
AB		PADDOCK LLC	60MG	A201291	001	May 24, 2013

TABLET;ORAL

TROSPIUM CHLORIDE

AB		APOTEX	20MG	A091513	001	Dec 06, 2011
AB	!	GLENMARK GENERICS	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		PADDOCK LLC	20MG	A091573	001	Nov 17, 2010

TRYPAN BLUE

SOLUTION;OPHTHALMIC

MEMBRANEBLUE

+	!	DORC	0.15%	N022278	001	Feb 20, 2009
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VISIONBLUE

+	!	DORC	0.06%	N021670	001	Dec 16, 2004
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UBROGEPANT

TABLET;ORAL

UBRELVY

+		ALLERGAN	50MG	N211765	001	Dec 23, 2019
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+	!		100MG	N211765	002	Dec 23, 2019
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ULIPRISTAL ACETATE

TABLET;ORAL

ELLA

AB	+	!	LAB HRA PHARMA	30MG	N022474	001	Aug 13, 2010
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LOGILTA

AB		TEVA PHARMS USA	30MG	A207952	001	Feb 13, 2017
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UMECLIDINIUM BROMIDE

POWDER; INHALATION

INCRUSE ELLIPTA

+	!	GLAXO GRP ENGLAND	EQ 62.5MCG BASE/INH	N205382	001	Apr 30, 2014
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UMECLIDINIUM BROMIDE; VILANTEROL TRIFENATATE

POWDER; INHALATION

ANORO ELLIPTA

+	!	GLAXOSMITHKLINE	EQ 0.0625MG BASE/INH;EQ 0.025MG	N203975	001	Dec 18, 2013
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BASE/INH

UPADACITINIB

TABLET, EXTENDED RELEASE;ORAL

RINVOQ

+	!	ABBVIE INC	15MG	N211675	001	Aug 16, 2019
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UREA, C-14

CAPSULE;ORAL

PYTEST

+	!	AVENT	1uCi	N020617	001	May 09, 1997
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PYTEST KIT

+	!	AVENT	1uCi	N020617	002	May 09, 1997
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URIDINE TRIACETATE

GRANULE;ORAL

VISTOGARD

+	!	WELLSTAT THERAP	10GM/PACKET	N208159	001	Dec 11, 2015
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XURIDEN

+	!	WELLSTAT THERAP	2GM/PACKET	N208169	001	Sep 04, 2015
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PRESCRIPTION DRUG PRODUCT LIST

URSODIOL

CAPSULE;ORAL

ACTIGALL

AB	+	ALLERGAN	300MG	N019594	002	Dec 31, 1987
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URSODIOL

AB		ABHAI LLC	300MG	A210707	001	May 17, 2018
AB		AMNEAL PHARMS CO	300MG	A211301	001	Oct 16, 2018
AB		EPIC PHARMA	300MG	A075517	001	Mar 14, 2000
AB		EYWA PHARMA	300MG	A212452	001	Oct 30, 2019
AB		LANNETT CO INC	300MG	A079082	001	Dec 15, 2008
AB		MYLAN	300MG	A090530	001	Feb 17, 2010
AB		RISING	300MG	A213200	001	Feb 12, 2020
AB		TEVA PHARMS	300MG	A075592	001	May 25, 2000

TABLET;ORAL

URSO 250

AB	+	ALLERGAN	250MG	N020675	001	Dec 10, 1997
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URSO FORTE

AB	+	ALLERGAN	500MG	N020675	002	Jul 21, 2004
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URSODIOL

AB		GLENMARK GENERICS	250MG	A090801	001	Jul 12, 2011
AB			500MG	A090801	002	Jul 12, 2011
AB		IMPAX LABS INC	250MG	A200826	001	Dec 23, 2011
AB			500MG	A200826	002	Dec 23, 2011
AB		PAR PHARM	250MG	A202540	001	Feb 14, 2013
AB			500MG	A202540	002	Feb 14, 2013
AB		ZYDUS	250MG	A211145	001	Oct 30, 2018
AB			500MG	A211145	002	Oct 30, 2018

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		CIPLA	EQ 500MG BASE	A077135	001	May 24, 2010
AB			EQ 1GM BASE	A077135	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		HIKMA	EQ 500MG BASE	A078656	001	May 24, 2010
AB			EQ 1GM BASE	A078656	002	May 24, 2010
AB		JUBILANT GENERICS	EQ 500MG BASE	A201506	001	Apr 03, 2012
AB			EQ 1GM BASE	A201506	002	Apr 03, 2012
AB		MYLAN PHARMS INC	EQ 500MG BASE	A078518	001	May 24, 2010
AB			EQ 1GM BASE	A078518	002	May 24, 2010
AB		SANDOZ	EQ 500MG BASE	A077478	001	May 24, 2010
AB			EQ 1GM BASE	A077478	002	May 24, 2010
AB		SUN PHARM INDS LTD	EQ 500MG BASE	A076588	001	Jan 31, 2007
AB			EQ 1GM BASE	A076588	002	Jan 31, 2007
AB		TIME-CAP LABS INC	EQ 500MG BASE	A079012	001	May 24, 2010
AB			EQ 1GM BASE	A079012	002	May 24, 2010
AB		WOCKHARDT	EQ 500MG BASE	A090216	001	May 24, 2010
AB			EQ 1GM BASE	A090216	002	May 24, 2010
AB		YILING PHARM LTD	EQ 500MG BASE	A209553	001	Mar 18, 2020
AB			EQ 1GM BASE	A209553	002	Mar 18, 2020
AB		ZYDUS PHARMS	EQ 500MG BASE	A079137	001	Dec 29, 2017
AB			EQ 1GM BASE	A079137	002	Dec 29, 2017

VALTREX

AB	+	GLAXOSMITHKLINE	EQ 500MG BASE	N020487	001	Jun 23, 1995
AB	+		EQ 1GM BASE	N020487	002	Jun 23, 1995

VALBENAZINE TOSYLATE

CAPSULE;ORAL

INGREZZA

	+	NEUROCRINE	EQ 40MG BASE	N209241	001	Apr 11, 2017
	+		EQ 80MG BASE	N209241	002	Oct 04, 2017

VALGANCICLOVIR HYDROCHLORIDE

FOR SOLUTION;ORAL

VALCYTE

AB	+	HOFFMANN LA ROCHE	50MG/ML	N022257	001	Aug 28, 2009
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VALGANCICLOVIR HYDROCHLORIDE

AB		ACTAVIS LABS FL INC	50MG/ML	A205220	001	Jul 18, 2016
AB		AJANTA PHARMA LTD	50MG/ML	A212890	001	Jan 13, 2020

PRESCRIPTION DRUG PRODUCT LIST

VALGANCICLOVIR HYDROCHLORIDE

FOR SOLUTION;ORAL

VALGANCICLOVIR HYDROCHLORIDE

AB GRANULES PHARMS **50MG/ML** **A213306 001** Jan 31, 2020
 TABLET;ORAL

VALCYTE

AB +! HOFFMANN LA ROCHE **EQ 450MG BASE** **N021304 001** Mar 29, 2001

VALGANCICLOVIR HYDROCHLORIDE

AB AJANTA PHARMA LTD **EQ 450MG BASE** **A212234 001** Dec 26, 2019
AB AUROBINDO PHARMA LTD **EQ 450MG BASE** **A204750 001** Mar 31, 2016
AB CIPLA **EQ 450MG BASE** **A209672 001** Nov 09, 2018
AB DR REDDYS **EQ 450MG BASE** **A206876 001** Dec 12, 2017
AB DR REDDYS LABS LTD **EQ 450MG BASE** **A203511 001** Nov 04, 2014
AB ENDO PHARMS INC **EQ 450MG BASE** **A200790 001** Nov 04, 2014
AB HETERO LABS LTD V **EQ 450MG BASE** **A205166 001** Mar 18, 2016

VALPROATE SODIUM

INJECTABLE;INJECTION

VALPROATE SODIUM

AP ATHENEX INC **EQ 100MG BASE/ML** **A076295 001** Nov 14, 2002
AP FRESENIUS KABI USA **EQ 100MG BASE/ML** **A076539 001** Jun 26, 2003
AP ! HIKMA FARMACEUTICA **EQ 100MG BASE/ML** **A078523 001** Feb 17, 2010

VALPROIC ACID

CAPSULE;ORAL

VALPROIC ACID

AB ! BIONPHARMA INC **250MG** **A073484 001** Jun 29, 1993
AB CATALENT **250MG** **A073229 001** Oct 29, 1991
AB NOVELGENIX THERAPS **250MG** **A207611 001** Aug 05, 2019
AB SUN PHARM INDS LTD **250MG** **A091037 001** Feb 22, 2013

SYRUP;ORAL

VALPROIC ACID

AA ECI PHARMS LLC **250MG/5ML** **A090517 001** May 28, 2010
AA HIGH TECH PHARMA **250MG/5ML** **A074060 001** Jan 13, 1995
AA LANNETT CO INC **250MG/5ML** **A077960 001** Oct 13, 2006
AA ! PHARM ASSOC **250MG/5ML** **A075379 001** Dec 15, 2000
AA VISTAPHARM **250MG/5ML** **A075782 001** Dec 22, 2000
AA WOCKHARDT BIO AG **250MG/5ML** **A070868 001** Jul 01, 1986

VALRUBICIN

SOLUTION;INTRAVESICAL

VALRUBICIN

AO CUSTOPHARM INC **40MG/ML** **A206430 001** Apr 19, 2019
VALSTAR PRESERVATIVE FREE
AO +! ENDO PHARM **40MG/ML** **N020892 001** Sep 25, 1998

VALSARTAN

TABLET;ORAL

DIOVAN

AB + NOVARTIS **40MG** **N021283 004** Aug 14, 2002
AB + **80MG** **N021283 001** Jul 18, 2001
AB + **160MG** **N021283 002** Jul 18, 2001
AB +! **320MG** **N021283 003** Jul 18, 2001

VALSARTAN

AB ALEMBIC PHARMS LTD **40MG** **A091367 001** Jan 05, 2015
AB **80MG** **A091367 002** Jan 05, 2015
AB **160MG** **A091367 003** Jan 05, 2015
AB **320MG** **A091367 004** Jan 05, 2015
AB ALKEM LABS LTD **40MG** **A205536 001** Mar 12, 2019
AB **80MG** **A205536 002** Mar 12, 2019
AB **160MG** **A205536 003** Mar 12, 2019
AB **320MG** **A205536 004** Mar 12, 2019
AB AMNEAL PHARMS **40MG** **A204011 001** Jan 11, 2016
AB **80MG** **A204011 002** Jan 11, 2016
AB **160MG** **A204011 003** Jan 11, 2016
AB **320MG** **A204011 004** Jan 11, 2016
AB AUROBINDO PHARMA LTD **40MG** **A202223 001** Jan 05, 2015
AB **80MG** **A202223 002** Jan 05, 2015
AB **160MG** **A202223 003** Jan 05, 2015
AB **320MG** **A202223 004** Jan 05, 2015
AB HETERO LABS LTD V **40MG** **A203311 001** Jan 05, 2015
AB **80MG** **A203311 002** Jan 05, 2015
AB **160MG** **A203311 003** Jan 05, 2015

PRESCRIPTION DRUG PRODUCT LIST

VALSARTAN

TABLET; ORAL

VALSARTAN

<u>AB</u>		<u>320MG</u>	<u>A203311 004</u>	Jan 05, 2015
<u>AB</u>	IVAX PHARMS	<u>40MG</u>	<u>A077530 001</u>	Jan 04, 2016
<u>AB</u>		<u>80MG</u>	<u>A077530 002</u>	Jan 04, 2016
<u>AB</u>		<u>160MG</u>	<u>A077530 003</u>	Jan 04, 2016
<u>AB</u>		<u>320MG</u>	<u>A077530 004</u>	Jan 04, 2016
<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>	SQUARE PHARMS LTD	<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>	UNICHEM LABS LTD	<u>40MG</u>	<u>A209261 001</u>	May 04, 2018
<u>AB</u>		<u>80MG</u>	<u>A209261 002</u>	May 04, 2018
<u>AB</u>		<u>160MG</u>	<u>A209261 003</u>	May 04, 2018
<u>AB</u>		<u>320MG</u>	<u>A209261 004</u>	May 04, 2018

VANCOMYCIN HYDROCHLORIDE

CAPSULE; ORAL

VANCOCIN HYDROCHLORIDE

<u>AB</u>	+	ANI PHARMS INC	<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
<u>VANCOMYCIN HYDROCHLORIDE</u>					
<u>AB</u>		AKORN	<u>EQ 125MG BASE</u>	<u>A065478 001</u>	Apr 09, 2012
<u>AB</u>			<u>EQ 250MG BASE</u>	<u>A065478 002</u>	Apr 09, 2012
<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
<u>AB</u>		WATSON LABS	<u>EQ 125MG BASE</u>	<u>A065510 001</u>	Apr 09, 2012
<u>AB</u>			<u>EQ 250MG BASE</u>	<u>A065510 002</u>	Apr 09, 2012

FOR SOLUTION; ORAL

FIRVANQ KIT

+

+

VANCOCIN HYDROCHLORIDE

ANI PHARMS INC

INJECTABLE; INJECTION

VANCOMYCIN HYDROCHLORIDE

<u>AP</u>		AUROBINDO PHARMA LTD	<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

PRESCRIPTION DRUG PRODUCT LIST

VANCOMYCIN HYDROCHLORIDE

INJECTABLE; INJECTION

VANCOMYCIN HYDROCHLORIDE

<u>AP</u>	!		<u>EQ 1GM BASE/VIAL</u>	<u>A062663</u>	<u>002</u>	Jul 31, 1987
<u>AP</u>	!		<u>EQ 5GM BASE/VIAL</u>	<u>A062663</u>	<u>003</u>	Jun 03, 1988
<u>AP</u>	!		<u>EQ 10GM BASE/VIAL</u>	<u>A062663</u>	<u>004</u>	Nov 28, 1997
<u>AP</u>		GLAND PHARMA LTD	<u>EQ 500MG BASE/VIAL</u>	<u>A205694</u>	<u>001</u>	Jan 21, 2016
<u>AP</u>			<u>EQ 1GM BASE/VIAL</u>	<u>A205694</u>	<u>002</u>	Jan 21, 2016
<u>AP</u>		HIKMA	<u>EQ 5GM BASE/VIAL</u>	<u>A204360</u>	<u>001</u>	Oct 15, 2018
<u>AP</u>			<u>EQ 10GM BASE/VIAL</u>	<u>A204360</u>	<u>002</u>	Oct 15, 2018
<u>AP</u>		HIKMA PHARMS	<u>EQ 750MG BASE/VIAL</u>	<u>A206616</u>	<u>001</u>	Oct 03, 2018
<u>AP</u>	!	HOSPIRA	<u>EQ 500MG BASE/VIAL</u>	<u>A062911</u>	<u>001</u>	Aug 04, 1988
<u>AP</u>	!		<u>EQ 500MG BASE/VIAL</u>	<u>A062931</u>	<u>001</u>	Oct 29, 1992
<u>AP</u>	!		<u>EQ 750MG BASE/VIAL</u>	<u>A062912</u>	<u>002</u>	Jan 07, 2009
<u>AP</u>	!		<u>EQ 750MG BASE/VIAL</u>	<u>A062933</u>	<u>002</u>	May 27, 2009
<u>AP</u>	!		<u>EQ 1GM BASE/VIAL</u>	<u>A062912</u>	<u>001</u>	Aug 04, 1988
<u>AP</u>	!		<u>EQ 1GM BASE/VIAL</u>	<u>A062933</u>	<u>001</u>	Oct 29, 1992
<u>AP</u>	!		<u>EQ 5GM BASE/VIAL</u>	<u>A063076</u>	<u>001</u>	Dec 21, 1990
<u>AP</u>		HOSPIRA INC	<u>EQ 10GM BASE/VIAL</u>	<u>A065455</u>	<u>001</u>	Apr 29, 2009
<u>AP</u>		MYLAN LABS LTD	<u>EQ 500MG BASE/VIAL</u>	<u>A065397</u>	<u>001</u>	Dec 30, 2008
<u>AP</u>			<u>EQ 1GM BASE/VIAL</u>	<u>A065397</u>	<u>002</u>	Dec 30, 2008
<u>AP</u>			<u>EQ 5GM BASE/VIAL</u>	<u>A065432</u>	<u>001</u>	Dec 30, 2008
<u>AP</u>			<u>EQ 10GM BASE/VIAL</u>	<u>A091554</u>	<u>001</u>	Sep 19, 2011
<u>AP</u>		PHARM ASSOC	<u>EQ 500MG BASE/VIAL</u>	<u>A065401</u>	<u>001</u>	Jun 30, 2008
<u>AP</u>			<u>EQ 1GM BASE/VIAL</u>	<u>A065401</u>	<u>002</u>	Jun 30, 2008
<u>AP</u>		SAGENT PHARMS	<u>EQ 5GM BASE/VIAL</u>	<u>A200837</u>	<u>001</u>	Aug 10, 2012
<u>AP</u>			<u>EQ 10GM BASE/VIAL</u>	<u>A200837</u>	<u>002</u>	Sep 02, 2014
<u>AP</u>		SANDOZ	<u>EQ 500MG BASE/VIAL</u>	<u>A090250</u>	<u>001</u>	Apr 27, 2010
<u>AP</u>			<u>EQ 1GM BASE/VIAL</u>	<u>A090250</u>	<u>002</u>	Apr 27, 2010
<u>AP</u>		SANDOZ INC	<u>EQ 5GM BASE/VIAL</u>	<u>A201048</u>	<u>001</u>	Aug 10, 2012
<u>AP</u>			<u>EQ 10GM BASE/VIAL</u>	<u>A201048</u>	<u>002</u>	Aug 10, 2012
<u>AP</u>		XELLIA PHARMS APS	<u>EQ 5GM BASE/VIAL</u>	<u>A204125</u>	<u>001</u>	Dec 28, 2015
<u>AP</u>			<u>EQ 10GM BASE/VIAL</u>	<u>A204125</u>	<u>002</u>	Dec 28, 2015
<u>AP</u>			<u>EQ 500MG BASE/VIAL</u>	<u>A204107</u>	<u>001</u>	Dec 28, 2015
<u>AP</u>			<u>EQ 1GM BASE/VIAL</u>	<u>A204107</u>	<u>002</u>	Dec 28, 2015

VANCOICIN HYDROCHLORIDE IN PLASTIC CONTAINER

+	!	BAXTER HLTHCARE	EQ 500MG BASE/100ML	N050671	001	Apr 29, 1993
+	!		EQ 750MG BASE/150ML	N050671	002	Dec 20, 2010
+	!		EQ 1GM BASE/200ML	N050671	003	Mar 01, 1999

POWDER; INTRAVENOUS

VANCOMYCIN HYDROCHLORIDE

+	!	MYLAN LABS LTD	EQ 250MG BASE/VIAL	N209481	001	Jul 10, 2018
+	!		EQ 750MG BASE/VIAL	N209481	002	Jul 10, 2018
+	!		EQ 1.25GM BASE/VIAL	N209481	003	Jul 10, 2018
+	!		EQ 1.5GM BASE/VIAL	N209481	004	Jul 10, 2018

VANCOMYCIN HYDROCHLORIDE IN PLASTIC CONTAINER

		SAMSON MEDCL	EQ 100GM BASE	A091532	001	Jan 06, 2016
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SOLUTION; INTRAVENOUS

VANCOMYCIN HYDROCHLORIDE

+	!	XELLIA PHARMS APS	EQ 500MG BASE/100ML	N211962	001	Feb 15, 2019
+	!		EQ 1GM BASE/200ML	N211962	002	Feb 15, 2019
+	!		EQ 1.5GM BASE/300ML	N211962	003	Feb 15, 2019
+	!		EQ 2GM BASE/400ML	N211962	004	Feb 15, 2019

VANDETANIB

TABLET; ORAL

CAPRELSA

+		GENZYME CORP	100MG	N022405	001	Apr 06, 2011
+	!		300MG	N022405	002	Apr 06, 2011

VARDENAFIL HYDROCHLORIDE

TABLET; ORAL

LEVITRA

<u>AB</u>	+	BAYER HLTHCARE	<u>5MG</u>	<u>N021400</u>	<u>001</u>	Aug 19, 2003
<u>AB</u>	+		<u>10MG</u>	<u>N021400</u>	<u>002</u>	Aug 19, 2003
<u>AB</u>	+		<u>20MG</u>	<u>N021400</u>	<u>004</u>	Aug 19, 2003

VARDENAFIL HYDROCHLORIDE

<u>AB</u>		CROSSMEDIKA SA	<u>2.5MG</u>	<u>A209057</u>	<u>001</u>	Oct 31, 2018
<u>AB</u>			<u>5MG</u>	<u>A209057</u>	<u>002</u>	Oct 31, 2018
<u>AB</u>			<u>10MG</u>	<u>A209057</u>	<u>003</u>	Oct 31, 2018
<u>AB</u>			<u>20MG</u>	<u>A209057</u>	<u>004</u>	Oct 31, 2018
<u>AB</u>		MACLEODS PHARMS LTD	<u>2.5MG</u>	<u>A204632</u>	<u>001</u>	Oct 22, 2019
<u>AB</u>			<u>5MG</u>	<u>A204632</u>	<u>002</u>	Oct 22, 2019

PRESCRIPTION DRUG PRODUCT LIST

VARDENAFIL HYDROCHLORIDE

TABLET; ORAL

VARDENAFIL HYDROCHLORIDE

<u>AB</u>		<u>10MG</u>	<u>A204632 003</u>	Oct 22, 2019
<u>AB</u>		<u>20MG</u>	<u>A204632 004</u>	Oct 22, 2019
<u>AB</u>	TEVA PHARMS	<u>2.5MG</u>	<u>A091347 001</u>	May 03, 2012
<u>AB</u>		<u>5MG</u>	<u>A091347 002</u>	May 03, 2012
<u>AB</u>		<u>10MG</u>	<u>A091347 003</u>	May 03, 2012
<u>AB</u>		<u>20MG</u>	<u>A091347 004</u>	May 03, 2012
<u>AB</u>	ZYDUS PHARMS	<u>2.5MG</u>	<u>A208960 001</u>	Oct 31, 2018
<u>AB</u>		<u>5MG</u>	<u>A208960 002</u>	Oct 31, 2018
<u>AB</u>		<u>10MG</u>	<u>A208960 003</u>	Oct 31, 2018
<u>AB</u>		<u>20MG</u>	<u>A208960 004</u>	Oct 31, 2018

TABLET, ORALLY DISINTEGRATING; ORAL

STAXYN

<u>AB</u>	+!	BAYER HLTHCARE	<u>10MG</u>	<u>N200179 001</u>	Jun 17, 2010
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VARDENAFIL HYDROCHLORIDE

<u>AB</u>		ALEMBIC PHARMS LTD	<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

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

+	!	PAR STERILE PRODUCTS	20UNITS/ML (20UNITS/ML)	N204485 001	Apr 17, 2014
+	!		200UNITS/10ML (20UNITS/ML)	N204485 002	Dec 17, 2016

VECURONIUM BROMIDE

INJECTABLE; INJECTION

VECURONIUM BROMIDE

<u>AP</u>		AUROBINDO PHARMA LTD	<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 PHARMA LTD	<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>A203725 001</u>	Jul 30, 2019
<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>		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
<u>AP</u>		TEVA PHARMS USA	<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>		WEST-WARD PHARMS INT	<u>10MG/VIAL</u>	<u>A075549 001</u>	Jun 13, 2000
<u>AP</u>			<u>20MG/VIAL</u>	<u>A075549 002</u>	Jun 13, 2000

VELAGLUCERASE ALFA

INJECTABLE; INTRAVENOUS

VPRIV

	SHIRE HUMAN GENETIC	400 UNITS/VIAL	N022575 001	Feb 26, 2010
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VEMURAFENIB

TABLET; ORAL

ZELBORAF

+	!	HOFFMANN LA ROCHE	240MG	N202429 001	Aug 17, 2011
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VENETOCLAX

TABLET; ORAL

VENCLEXTA

+		ABBVIE INC	10MG	N208573 001	Apr 11, 2016
+			50MG	N208573 002	Apr 11, 2016
+	!		100MG	N208573 003	Apr 11, 2016

PRESCRIPTION DRUG PRODUCT LIST

VENLAFAXINE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE;ORAL

EFFEXOR XR

<u>AB</u>	<u>+</u>	<u>WYETH PHARMS</u>	<u>EQ 37.5MG BASE</u>	<u>N020699 001</u>	<u>Oct 20, 1997</u>
<u>AB</u>	<u>+</u>		<u>EQ 75MG BASE</u>	<u>N020699 002</u>	<u>Oct 20, 1997</u>
<u>AB</u>	<u>+</u>		<u>EQ 150MG BASE</u>	<u>N020699 004</u>	<u>Oct 20, 1997</u>

VENLAFAXINE HYDROCHLORIDE

<u>AB</u>		<u>ANNORA PHARMA</u>	<u>EQ 37.5MG BASE</u>	<u>A212277 001</u>	<u>Jul 08, 2019</u>
<u>AB</u>			<u>EQ 75MG BASE</u>	<u>A212277 002</u>	<u>Jul 08, 2019</u>
<u>AB</u>			<u>EQ 150MG BASE</u>	<u>A212277 003</u>	<u>Jul 08, 2019</u>
<u>AB</u>		<u>AUROBINDO PHARMA LTD</u>	<u>EQ 37.5MG BASE</u>	<u>A200834 001</u>	<u>Apr 14, 2011</u>
<u>AB</u>			<u>EQ 75MG BASE</u>	<u>A200834 002</u>	<u>Apr 14, 2011</u>
<u>AB</u>			<u>EQ 150MG BASE</u>	<u>A200834 003</u>	<u>Apr 14, 2011</u>
<u>AB</u>		<u>DR REDDYS LABS LTD</u>	<u>EQ 37.5MG BASE</u>	<u>A078421 001</u>	<u>May 06, 2011</u>
<u>AB</u>			<u>EQ 75MG BASE</u>	<u>A078421 002</u>	<u>May 06, 2011</u>
<u>AB</u>			<u>EQ 150MG BASE</u>	<u>A078421 003</u>	<u>May 06, 2011</u>
<u>AB</u>		<u>INTELLIPHARMACEUTICS</u>	<u>EQ 37.5MG BASE</u>	<u>A201272 001</u>	<u>Nov 23, 2018</u>
<u>AB</u>			<u>EQ 75MG BASE</u>	<u>A201272 002</u>	<u>Nov 23, 2018</u>
<u>AB</u>			<u>EQ 150MG BASE</u>	<u>A201272 003</u>	<u>Nov 23, 2018</u>
<u>AB</u>		<u>INVENTIA HLTHCARE</u>	<u>EQ 37.5MG BASE</u>	<u>A203332 001</u>	<u>Mar 12, 2020</u>
<u>AB</u>			<u>EQ 75MG BASE</u>	<u>A203332 002</u>	<u>Mar 12, 2020</u>
<u>AB</u>			<u>EQ 150MG BASE</u>	<u>A203332 003</u>	<u>Mar 12, 2020</u>
<u>AB</u>		<u>MACLEODS PHARMS LTD</u>	<u>EQ 37.5MG BASE</u>	<u>A204889 001</u>	<u>Oct 05, 2017</u>
<u>AB</u>			<u>EQ 75MG BASE</u>	<u>A204889 002</u>	<u>Oct 05, 2017</u>
<u>AB</u>			<u>EQ 150MG BASE</u>	<u>A204889 003</u>	<u>Oct 05, 2017</u>
<u>AB</u>		<u>ORCHID HLTHCARE</u>	<u>EQ 37.5MG BASE</u>	<u>A091123 001</u>	<u>Jul 11, 2011</u>
<u>AB</u>			<u>EQ 75MG BASE</u>	<u>A091123 002</u>	<u>Jul 11, 2011</u>
<u>AB</u>			<u>EQ 150MG BASE</u>	<u>A091123 003</u>	<u>Jul 11, 2011</u>
<u>AB</u>		<u>TEVA</u>	<u>EQ 37.5MG BASE</u>	<u>A076565 001</u>	<u>Jun 28, 2010</u>
<u>AB</u>			<u>EQ 75MG BASE</u>	<u>A076565 002</u>	<u>Jun 28, 2010</u>
<u>AB</u>			<u>EQ 150MG BASE</u>	<u>A076565 003</u>	<u>Jun 28, 2010</u>
<u>AB</u>		<u>TORRENT PHARMS LLC</u>	<u>EQ 37.5MG BASE</u>	<u>A090899 001</u>	<u>Jun 01, 2011</u>
<u>AB</u>			<u>EQ 75MG BASE</u>	<u>A090899 002</u>	<u>Jun 01, 2011</u>
<u>AB</u>			<u>EQ 150MG BASE</u>	<u>A090899 003</u>	<u>Jun 01, 2011</u>
<u>AB</u>		<u>VALEANT PHARMS NORTH</u>	<u>EQ 37.5MG BASE</u>	<u>A090071 001</u>	<u>Apr 15, 2011</u>
<u>AB</u>			<u>EQ 75MG BASE</u>	<u>A090071 002</u>	<u>Apr 15, 2011</u>
<u>AB</u>			<u>EQ 150MG BASE</u>	<u>A090071 003</u>	<u>Apr 15, 2011</u>
<u>AB</u>		<u>WOCKHARDT</u>	<u>EQ 37.5MG BASE</u>	<u>A078865 001</u>	<u>Apr 14, 2011</u>
<u>AB</u>			<u>EQ 75MG BASE</u>	<u>A078865 002</u>	<u>Apr 14, 2011</u>
<u>AB</u>			<u>EQ 150MG BASE</u>	<u>A078865 003</u>	<u>Apr 14, 2011</u>
<u>AB</u>		<u>ZYDUS PHARMS USA INC</u>	<u>EQ 37.5MG BASE</u>	<u>A090174 001</u>	<u>Apr 14, 2011</u>
<u>AB</u>			<u>EQ 75MG BASE</u>	<u>A090174 002</u>	<u>Apr 14, 2011</u>
<u>AB</u>			<u>EQ 150MG BASE</u>	<u>A090174 003</u>	<u>Apr 14, 2011</u>

TABLET;ORAL

VENLAFAXINE HYDROCHLORIDE

<u>AB</u>		<u>ALEMBIC PHARMS LTD</u>	<u>EQ 25MG BASE</u>	<u>A078932 001</u>	<u>Dec 14, 2010</u>
<u>AB</u>			<u>EQ 37.5MG BASE</u>	<u>A078932 002</u>	<u>Dec 14, 2010</u>
<u>AB</u>			<u>EQ 50MG BASE</u>	<u>A078932 003</u>	<u>Dec 14, 2010</u>
<u>AB</u>			<u>EQ 75MG BASE</u>	<u>A078932 004</u>	<u>Dec 14, 2010</u>
<u>AB</u>			<u>EQ 100MG BASE</u>	<u>A078932 005</u>	<u>Dec 14, 2010</u>
<u>AB</u>		<u>AMNEAL PHARMS</u>	<u>EQ 25MG BASE</u>	<u>A079098 001</u>	<u>May 11, 2010</u>
<u>AB</u>			<u>EQ 37.5MG BASE</u>	<u>A079098 002</u>	<u>May 11, 2010</u>
<u>AB</u>			<u>EQ 50MG BASE</u>	<u>A079098 003</u>	<u>May 11, 2010</u>
<u>AB</u>			<u>EQ 75MG BASE</u>	<u>A079098 004</u>	<u>May 11, 2010</u>
<u>AB</u>			<u>EQ 100MG BASE</u>	<u>A079098 005</u>	<u>May 11, 2010</u>
<u>AB</u>		<u>AUROBINDO PHARMA</u>	<u>EQ 25MG BASE</u>	<u>A090555 001</u>	<u>Apr 07, 2010</u>
<u>AB</u>			<u>EQ 37.5MG BASE</u>	<u>A090555 002</u>	<u>Apr 07, 2010</u>
<u>AB</u>			<u>EQ 50MG BASE</u>	<u>A090555 003</u>	<u>Apr 07, 2010</u>
<u>AB</u>			<u>EQ 75MG BASE</u>	<u>A090555 004</u>	<u>Apr 07, 2010</u>
<u>AB</u>			<u>EQ 100MG BASE</u>	<u>A090555 005</u>	<u>Apr 07, 2010</u>
<u>AB</u>		<u>CADILA PHARMS LTD</u>	<u>EQ 25MG BASE</u>	<u>A206250 001</u>	<u>Nov 21, 2018</u>
<u>AB</u>			<u>EQ 37.5MG BASE</u>	<u>A206250 002</u>	<u>Nov 21, 2018</u>
<u>AB</u>			<u>EQ 50MG BASE</u>	<u>A206250 003</u>	<u>Nov 21, 2018</u>
<u>AB</u>			<u>EQ 75MG BASE</u>	<u>A206250 004</u>	<u>Nov 21, 2018</u>
<u>AB</u>			<u>EQ 100MG BASE</u>	<u>A206250 005</u>	<u>Nov 21, 2018</u>
<u>AB</u>		<u>DR REDDYS LABS LTD</u>	<u>EQ 25MG BASE</u>	<u>A078301 001</u>	<u>Jun 13, 2008</u>
<u>AB</u>			<u>EQ 37.5MG BASE</u>	<u>A078301 002</u>	<u>Jun 13, 2008</u>
<u>AB</u>			<u>EQ 50MG BASE</u>	<u>A078301 003</u>	<u>Jun 13, 2008</u>
<u>AB</u>			<u>EQ 75MG BASE</u>	<u>A078301 004</u>	<u>Jun 13, 2008</u>

PRESCRIPTION DRUG PRODUCT LIST

3-444 (of 453)

VENLAFAXINE HYDROCHLORIDE

TABLET; ORAL

VENLAFAXINE HYDROCHLORIDE

<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A078301 005</u>	Jun 13, 2008
<u>AB</u>	HERITAGE PHARMS INC	<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>	MYLAN	<u>EQ 100MG BASE</u>	<u>A078554 005</u>	Jan 09, 2009
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>A077166 001</u>	Jun 13, 2008
<u>AB</u>		<u>EQ 37.5MG BASE</u>	<u>A077166 002</u>	Jun 13, 2008
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A077166 003</u>	Jun 13, 2008
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A077166 004</u>	Jun 13, 2008
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A077166 005</u>	Jun 13, 2008
<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
<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>	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 PHARMA	<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>	+ OSMOTICA PHARM	<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>	SUN PHARM	<u>EQ 37.5MG BASE</u>	<u>A091272 001</u>	Aug 18, 2010
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A091272 002</u>	Aug 18, 2010
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A091272 003</u>	Aug 18, 2010
<u>AB</u>		<u>EQ 225MG BASE</u>	<u>A091272 004</u>	Jan 08, 2019

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

VERELAN

<u>AB</u>	+ RECRO GAINESVILLE	<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>AB</u>	+!	<u>360MG</u>	<u>N019614 004</u>	May 10, 1996
	VERELAN PM			
	+ RECRO GAINESVILLE	<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

SOLUTION; INTRAVENOUS

VERAPAMIL HYDROCHLORIDE

<u>AP</u>	AMNEAL	<u>5MG/2ML (2.5MG/ML)</u>	<u>A210994 001</u>	Jul 13, 2018
<u>AP</u>		<u>10MG/4ML (2.5MG/ML)</u>	<u>A210994 002</u>	Jul 13, 2018
<u>AP</u>	AREVA PHARMS	<u>10MG/4ML (2.5MG/ML)</u>	<u>A213352 001</u>	Mar 17, 2020
<u>AP</u>	EXELA PHARMA SCS	<u>5MG/2ML (2.5MG/ML)</u>	<u>N018925 001</u>	Mar 30, 1984

PRESCRIPTION DRUG PRODUCT LIST

VERAPAMIL HYDROCHLORIDE

SOLUTION; INTRAVENOUS

VERAPAMIL HYDROCHLORIDE

LLC

<u>AP</u>		<u>10MG/4ML (2.5MG/ML)</u>	<u>N018925 002</u>	Apr 05, 2018
<u>AP</u>	!	<u>5MG/2ML (2.5MG/ML)</u>	<u>A070738 001</u>	May 06, 1987
<u>AP</u>	!	<u>5MG/2ML (2.5MG/ML)</u>	<u>A075136 001</u>	Oct 20, 1998
<u>AP</u>	!	<u>5MG/2ML (2.5MG/ML)</u>	<u>A070737 001</u>	May 06, 1987
<u>AP</u>	!	<u>10MG/4ML (2.5MG/ML)</u>	<u>A070737 002</u>	May 06, 1987
<u>AP</u>	MICRO LABS	<u>5MG/2ML (2.5MG/ML)</u>	<u>A211370 001</u>	Dec 28, 2018
<u>AP</u>		<u>10MG/4ML (2.5MG/ML)</u>	<u>A211370 002</u>	Dec 28, 2018
<u>AP</u>	SOMERSET	<u>5MG/2ML (2.5MG/ML)</u>	<u>A211035 001</u>	Jun 18, 2018
<u>AP</u>		<u>10MG/4ML (2.5MG/ML)</u>	<u>A211035 002</u>	Jun 18, 2018
<u>AP</u>	SOMERSET THERAPS LLC	<u>5MG/2ML (2.5MG/ML)</u>	<u>A211015 001</u>	Jun 18, 2018
<u>AP</u>		<u>10MG/4ML (2.5MG/ML)</u>	<u>A211015 002</u>	Jun 18, 2018

TABLET; ORAL

VERAPAMIL HYDROCHLORIDE

<u>AB</u>	HERITAGE PHARMS INC	<u>40MG</u>	<u>A071881 002</u>	Oct 14, 2015
<u>AB</u>		<u>80MG</u>	<u>A071881 003</u>	Apr 05, 1988
<u>AB</u>	!	<u>120MG</u>	<u>A071881 001</u>	Apr 05, 1988
<u>AB</u>	MYLAN	<u>80MG</u>	<u>A071483 002</u>	Feb 15, 1989
<u>AB</u>		<u>120MG</u>	<u>A071483 001</u>	Feb 15, 1989
<u>AB</u>	WATSON LABS	<u>40MG</u>	<u>A072924 001</u>	Jun 29, 1993
<u>AB</u>		<u>80MG</u>	<u>A070995 001</u>	Oct 01, 1986
<u>AB</u>		<u>120MG</u>	<u>A070994 001</u>	Oct 01, 1986

TABLET, EXTENDED RELEASE; ORAL

CALAN SR

<u>AB</u>	+	PFIZER	<u>120MG</u>	<u>N019152 003</u>	Mar 06, 1991
<u>AB</u>	+		<u>240MG</u>	<u>N019152 001</u>	Dec 16, 1986

VERAPAMIL HYDROCHLORIDE

<u>AB</u>		CADILA PHARMS LTD	<u>180MG</u>	<u>A206173 001</u>	May 05, 2017
<u>AB</u>			<u>240MG</u>	<u>A206173 002</u>	May 05, 2017
<u>AB</u>		GLENMARK GENERICS	<u>120MG</u>	<u>A090700 001</u>	Aug 03, 2011
<u>AB</u>	!		<u>180MG</u>	<u>A090700 002</u>	Aug 03, 2011
<u>AB</u>			<u>240MG</u>	<u>A078906 001</u>	Sep 17, 2009
<u>AB</u>		IVAX SUB TEVA PHARMS	<u>120MG</u>	<u>A073568 002</u>	Oct 10, 1997
<u>AB</u>			<u>180MG</u>	<u>A074330 001</u>	Jan 31, 1994
<u>AB</u>			<u>240MG</u>	<u>A073568 001</u>	Jul 31, 1992
<u>AB</u>		MYLAN	<u>120MG</u>	<u>A074587 002</u>	Feb 21, 1997
<u>AB</u>			<u>180MG</u>	<u>A074587 003</u>	Sep 09, 1997
<u>AB</u>			<u>240MG</u>	<u>A074587 001</u>	Mar 23, 1996
<u>AB</u>		PAR PHARM	<u>120MG</u>	<u>A075072 001</u>	May 25, 1999
<u>AB</u>			<u>240MG</u>	<u>A075072 003</u>	May 25, 1999
<u>AB</u>		SUN PHARM INDS INC	<u>120MG</u>	<u>A090529 001</u>	Dec 30, 2011
<u>AB</u>			<u>180MG</u>	<u>A090529 002</u>	Dec 30, 2011
<u>AB</u>			<u>240MG</u>	<u>A090529 003</u>	Dec 30, 2011

VERTEPORFIN

INJECTABLE; INJECTION

VISUDYNE

+	VALEANT LUXEMBOURG	15MG/VIAL	<u>N021119 001</u>	Apr 12, 2000
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VIGABATRIN

FOR SOLUTION; ORAL

SABRIL

<u>AA</u>	+	LUNDBECK PHARMS LLC	<u>500MG/PACKET</u>	<u>N022006 001</u>	Aug 21, 2009
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VIGABATRIN

<u>AA</u>		AMNEAL PHARMS	<u>500MG/PACKET</u>	<u>A210155 001</u>	Mar 13, 2018
<u>AA</u>		DR REDDYS LABS LTD	<u>500MG/PACKET</u>	<u>A211481 001</u>	Nov 20, 2018
<u>AA</u>		INVAGEN PHARMS	<u>500MG/PACKET</u>	<u>A211592 001</u>	Dec 03, 2019
<u>AA</u>		PAR PHARM INC	<u>500MG/PACKET</u>	<u>A208218 001</u>	Apr 27, 2017
<u>AA</u>		TEVA PHARMS USA	<u>500MG/PACKET</u>	<u>A209824 001</u>	Apr 23, 2018

VIGADRONE

<u>AA</u>		AUCTA	<u>500MG/PACKET</u>	<u>A210196 001</u>	Jun 21, 2018
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TABLET; ORAL

SABRIL

<u>AB</u>	+	LUNDBECK PHARMS LLC	<u>500MG</u>	<u>N020427 001</u>	Aug 21, 2009
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VIGABATRIN

<u>AB</u>		TEVA PHARMS USA	<u>500MG</u>	<u>A209822 001</u>	Jan 14, 2019
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PRESCRIPTION DRUG PRODUCT LIST

VILAZODONE HYDROCHLORIDE

TABLET; ORAL

VIIBRYD

AB	+ !	ALLERGAN	10MG	N022567 001	Jan 21, 2011
AB	+		20MG	N022567 002	Jan 21, 2011
AB	+		40MG	N022567 003	Jan 21, 2011

VILAZODONE HYDROCHLORIDE

AB		ALEMBIC PHARMS LTD	10MG	A208202 001	Jan 10, 2020
AB			20MG	A208202 002	Jan 10, 2020
AB			40MG	A208202 003	Jan 10, 2020
AB		TEVA PHARMS USA	10MG	A208212 001	Sep 30, 2019
AB			20MG	A208212 002	Sep 30, 2019
AB			40MG	A208212 003	Sep 30, 2019

VINBLASTINE SULFATE

INJECTABLE; INJECTION

VINBLASTINE SULFATE

!		FRESENIUS KABI USA	1MG/ML	A089515 001	Apr 29, 1987
!		WEST-WARD PHARMS INT	10MG/VIAL	A089395 001	Apr 09, 1987

VINCRISTINE SULFATE

INJECTABLE; INJECTION

VINCRISTINE SULFATE PFS

AP	!	HOSPIRA	1MG/ML	A071484 001	Apr 19, 1988
AP		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

VINORELBINE TARTRATE

AP		ACTAVIS TOTOWA	EQ 10MG BASE/ML	A078011 001	Jul 22, 2009
AP		DR REDDYS	EQ 10MG BASE/ML	A202017 001	Sep 12, 2013
AP		FRESENIUS KABI USA	EQ 10MG BASE/ML	A076849 001	Apr 18, 2005
AP		JIANGSU HANSOH PHARM	EQ 10MG BASE/ML	A091106 001	Sep 26, 2012
AP		TEVA PHARMS USA	EQ 10MG BASE/ML	A076028 001	Feb 03, 2003
AP		WEST-WARD PHARMS INT	EQ 10MG BASE/ML	A075992 001	Jun 10, 2003
AP			EQ 10MG BASE/ML	A076461 001	Dec 11, 2003

VISMODEGIB

CAPSULE; ORAL

ERIVEDGE

+ !		GENENTECH	150MG	N203388 001	Jan 30, 2012
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VITAMIN A PALMITATE

INJECTABLE; INJECTION

AQUASOL A

+ !		CASPER PHARMA LLC	EQ 50,000 UNITS BASE/ML	N006823 001	
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VORAPAXAR SULFATE

TABLET; ORAL

ZONTIVITY

+ !		TOPROL	EQ 2.08MG BASE	N204886 001	May 08, 2014
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VORICONAZOLE

FOR SUSPENSION; ORAL

VFEND

AB	+ !	PF PRISM CV	200MG/5ML	N021630 001	Dec 19, 2003
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VORICONAZOLE

AB		AMNEAL PHARMS	200MG/5ML	A205034 001	Apr 13, 2016
AB		NOVEL LABS INC	200MG/5ML	A206799 001	May 31, 2016

INJECTABLE; INTRAVENOUS

VFEND

AP	+ !	PF PRISM CV	200MG/VIAL	N021267 001	May 24, 2002
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VORICONAZOLE

AP		ALVOGEN	200MG/VIAL	A206398 001	Mar 23, 2016
AP		HAINAN POLY PHARM	200MG/VIAL	A211661 001	Nov 30, 2018
AP		SANDOZ INC	200MG/VIAL	A090862 001	May 30, 2012
AP		ZYDUS PHARMS	200MG/VIAL	A208983 001	Jul 16, 2018

POWDER; INTRAVENOUS

VORICONAZOLE

AP		XELLIA PHARMS APS	200MG/VIAL	N208562 001	Mar 09, 2017
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PRESCRIPTION DRUG PRODUCT LIST

VORICONAZOLE

TABLET; ORAL

VFEND

<u>AB</u>	+	PF PRISM CV	<u>50MG</u>	<u>N021266</u>	<u>001</u>	May 24, 2002
<u>AB</u>	+	!	<u>200MG</u>	<u>N021266</u>	<u>002</u>	May 24, 2002

VORICONAZOLE

<u>AB</u>		AJANTA PHARMA LTD	<u>50MG</u>	<u>A206181</u>	<u>001</u>	May 24, 2016
<u>AB</u>			<u>200MG</u>	<u>A206181</u>	<u>002</u>	May 24, 2016
<u>AB</u>		AKORN	<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>		APPCO	<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>		AUROBINDO PHARMA LTD	<u>50MG</u>	<u>A206837</u>	<u>001</u>	Jan 22, 2016
<u>AB</u>			<u>200MG</u>	<u>A206837</u>	<u>002</u>	Jan 22, 2016
<u>AB</u>		CADILA	<u>50MG</u>	<u>A206747</u>	<u>001</u>	May 24, 2016
<u>AB</u>			<u>200MG</u>	<u>A206747</u>	<u>002</u>	May 24, 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>		NOVEL LABS INC	<u>50MG</u>	<u>A207371</u>	<u>001</u>	May 24, 2016
<u>AB</u>			<u>200MG</u>	<u>A207371</u>	<u>002</u>	May 24, 2016
<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>		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

+	!	MERCK	100MG	N021991	001	Oct 06, 2006
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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

VOXELOTOR

TABLET; ORAL

OXBRYTA

+	!	GLOBAL BLOOD THERAPS	500MG	N213137	001	Nov 25, 2019
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WARFARIN SODIUM

TABLET; ORAL

COUMADIN

<u>AB</u>	+	BRISTOL MYERS SQUIBB	<u>1MG</u>	<u>N009218</u>	<u>022</u>	Mar 01, 1990
<u>AB</u>	+		<u>2MG</u>	<u>N009218</u>	<u>013</u>	
<u>AB</u>	+		<u>2.5MG</u>	<u>N009218</u>	<u>018</u>	
<u>AB</u>	+		<u>3MG</u>	<u>N009218</u>	<u>025</u>	Nov 18, 1996
<u>AB</u>	+		<u>4MG</u>	<u>N009218</u>	<u>023</u>	Aug 24, 1993
<u>AB</u>	+		<u>5MG</u>	<u>N009218</u>	<u>007</u>	
<u>AB</u>	+		<u>6MG</u>	<u>N009218</u>	<u>026</u>	Nov 18, 1996
<u>AB</u>	+		<u>7.5MG</u>	<u>N009218</u>	<u>016</u>	
<u>AB</u>	+	!	<u>10MG</u>	<u>N009218</u>	<u>005</u>	

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

PRESCRIPTION DRUG PRODUCT LIST

WARFARIN SODIUM

TABLET; ORAL

WARFARIN SODIUM

<u>AB</u>		<u>5MG</u>	<u>A202202 006</u>	Mar 04, 2013
<u>AB</u>		<u>6MG</u>	<u>A202202 007</u>	Mar 04, 2013
<u>AB</u>		<u>7.5MG</u>	<u>A202202 008</u>	Mar 04, 2013
<u>AB</u>		<u>10MG</u>	<u>A202202 009</u>	Mar 04, 2013
<u>AB</u>	BARR	<u>1MG</u>	<u>A040145 001</u>	Mar 26, 1997
<u>AB</u>		<u>2MG</u>	<u>A040145 002</u>	Mar 26, 1997
<u>AB</u>		<u>2.5MG</u>	<u>A040145 003</u>	Mar 26, 1997
<u>AB</u>		<u>3MG</u>	<u>A040145 008</u>	Nov 05, 1998
<u>AB</u>		<u>4MG</u>	<u>A040145 004</u>	Mar 26, 1997
<u>AB</u>		<u>5MG</u>	<u>A040145 005</u>	Mar 26, 1997
<u>AB</u>		<u>6MG</u>	<u>A040145 009</u>	Nov 05, 1998
<u>AB</u>		<u>7.5MG</u>	<u>A040145 006</u>	Mar 26, 1997
<u>AB</u>		<u>10MG</u>	<u>A040145 007</u>	Mar 26, 1997
<u>AB</u>	INVAGEN PHARMS	<u>1MG</u>	<u>A090935 001</u>	May 25, 2011
<u>AB</u>		<u>2MG</u>	<u>A090935 002</u>	May 25, 2011
<u>AB</u>		<u>2.5MG</u>	<u>A090935 003</u>	May 25, 2011
<u>AB</u>		<u>3MG</u>	<u>A090935 004</u>	May 25, 2011
<u>AB</u>		<u>4MG</u>	<u>A090935 005</u>	May 25, 2011
<u>AB</u>		<u>5MG</u>	<u>A090935 006</u>	May 25, 2011
<u>AB</u>		<u>6MG</u>	<u>A090935 007</u>	May 25, 2011
<u>AB</u>		<u>7.5MG</u>	<u>A090935 008</u>	May 25, 2011
<u>AB</u>		<u>10MG</u>	<u>A090935 009</u>	May 25, 2011
<u>AB</u>	IPCA LABS LTD	<u>1MG</u>	<u>A200104 001</u>	Jun 27, 2013
<u>AB</u>		<u>2MG</u>	<u>A200104 002</u>	Jun 27, 2013
<u>AB</u>		<u>2.5MG</u>	<u>A200104 003</u>	Jun 27, 2013
<u>AB</u>		<u>3MG</u>	<u>A200104 004</u>	Jun 27, 2013
<u>AB</u>		<u>4MG</u>	<u>A200104 005</u>	Jun 27, 2013
<u>AB</u>		<u>5MG</u>	<u>A200104 006</u>	Jun 27, 2013
<u>AB</u>		<u>6MG</u>	<u>A200104 007</u>	Jun 27, 2013
<u>AB</u>		<u>7.5MG</u>	<u>A200104 008</u>	Jun 27, 2013
<u>AB</u>		<u>10MG</u>	<u>A200104 009</u>	Jun 27, 2013
<u>AB</u>	PLIVA	<u>1MG</u>	<u>A040616 009</u>	Jul 05, 2006
<u>AB</u>		<u>2MG</u>	<u>A040616 001</u>	Jul 05, 2006
<u>AB</u>		<u>2.5MG</u>	<u>A040616 002</u>	Jul 05, 2006
<u>AB</u>		<u>3MG</u>	<u>A040616 003</u>	Jul 05, 2006
<u>AB</u>		<u>4MG</u>	<u>A040616 004</u>	Jul 05, 2006
<u>AB</u>		<u>5MG</u>	<u>A040616 005</u>	Jul 05, 2006
<u>AB</u>		<u>6MG</u>	<u>A040616 006</u>	Jul 05, 2006
<u>AB</u>		<u>7.5MG</u>	<u>A040616 007</u>	Jul 05, 2006
<u>AB</u>		<u>10MG</u>	<u>A040616 008</u>	Jul 05, 2006
<u>AB</u>	TARO	<u>1MG</u>	<u>A040301 002</u>	Jul 15, 1999
<u>AB</u>		<u>2MG</u>	<u>A040301 003</u>	Jul 15, 1999
<u>AB</u>		<u>2.5MG</u>	<u>A040301 004</u>	Jul 15, 1999
<u>AB</u>		<u>3MG</u>	<u>A040301 005</u>	Jul 15, 1999
<u>AB</u>		<u>4MG</u>	<u>A040301 006</u>	Jul 15, 1999
<u>AB</u>		<u>5MG</u>	<u>A040301 007</u>	Jul 15, 1999
<u>AB</u>		<u>6MG</u>	<u>A040301 008</u>	Jul 15, 1999
<u>AB</u>		<u>7.5MG</u>	<u>A040301 009</u>	Jul 15, 1999
<u>AB</u>		<u>10MG</u>	<u>A040301 001</u>	Jul 15, 1999
<u>AB</u>	ZYDUS PHARMS USA	<u>1MG</u>	<u>A040663 001</u>	May 30, 2006
<u>AB</u>		<u>2MG</u>	<u>A040663 002</u>	May 30, 2006
<u>AB</u>		<u>2.5MG</u>	<u>A040663 003</u>	May 30, 2006
<u>AB</u>		<u>3MG</u>	<u>A040663 004</u>	May 30, 2006
<u>AB</u>		<u>4MG</u>	<u>A040663 005</u>	May 30, 2006
<u>AB</u>		<u>5MG</u>	<u>A040663 006</u>	May 30, 2006
<u>AB</u>		<u>6MG</u>	<u>A040663 007</u>	May 30, 2006
<u>AB</u>		<u>7.5MG</u>	<u>A040663 008</u>	May 30, 2006
<u>AB</u>		<u>10MG</u>	<u>A040663 009</u>	May 30, 2006

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	

PRESCRIPTION DRUG PRODUCT LIST

ZAFIRLUKAST

TABLET; ORAL

ACCOLATE

AB	+	PAR PHARM INC	10MG	N020547	003	Sep 17, 1999
AB	+	!	20MG	N020547	001	Sep 26, 1996

ZAFIRLUKAST

AB		DR REDDYS LABS LTD	10MG	A090372	001	Nov 18, 2010
AB			20MG	A090372	002	Nov 18, 2010

ZALEPLON

CAPSULE; ORAL

SONATA

AB	+	PFIZER	5MG	N020859	001	Aug 13, 1999
AB	+	!	10MG	N020859	002	Aug 13, 1999

ZALEPLON

AB		AUROBINDO PHARMA	5MG	A078829	001	Jun 06, 2008
AB			10MG	A078829	002	Jun 06, 2008
AB		CIPLA LTD	5MG	A077505	001	Jun 20, 2008
AB			10MG	A077505	002	Jun 20, 2008
AB		HIKMA	5MG	A077237	001	Jun 06, 2008
AB			10MG	A077237	002	Jun 06, 2008
AB		HIKMA PHARMS	5MG	A078147	001	Nov 25, 2008
AB			10MG	A078147	002	Nov 25, 2008
AB		ORCHID HLTHCARE	5MG	A090374	001	Sep 17, 2009
AB			10MG	A090374	002	Sep 17, 2009
AB		TEVA PHARMS	5MG	A077239	001	Jun 06, 2008
AB			10MG	A077239	002	Jun 06, 2008
AB		UNICHEM	5MG	A078989	001	Jun 06, 2008
AB			10MG	A078989	002	Jun 06, 2008

ZANAMIVIR

POWDER; INHALATION

RELENZA

+	!	GLAXOSMITHKLINE	5MG	N021036	001	Jul 26, 1999
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ZANUBRUTINIB

CAPSULE; ORAL

BRUKINSA

+	!	BEIGENE	80MG	N213217	001	Nov 14, 2019
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ZICONOTIDE ACETATE

INJECTABLE; INTRATHECAL

PRIALT

+	!	TERSERA THERAPS LLC	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

ZIDOVDINE

CAPSULE; ORAL

RETROVIR

AB	+	VIIV HLTHCARE	100MG	N019655	001	Mar 19, 1987
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ZIDOVDINE

AB		AUROBINDO PHARMA LTD	100MG	A078128	001	Mar 27, 2006
AB		CIPLA LTD	100MG	A078349	001	May 23, 2007

INJECTABLE; INJECTION

RETROVIR

AP	+	VIIV HLTHCARE	10MG/ML	N019951	001	Feb 02, 1990
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SYRUP; ORAL

RETROVIR

AA	+	VIIV HLTHCARE	50MG/5ML	N019910	001	Sep 28, 1989
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ZIDOVDINE

AA		AUROBINDO	50MG/5ML	A077268	001	Sep 19, 2005
AA		CIPLA LTD	50MG/5ML	A077981	001	Jun 26, 2008

TABLET; ORAL

ZIDOVDINE

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, 2008
AB		HIKMA	300MG	A076844	001	Sep 19, 2005
AB		MYLAN PHARMS INC	300MG	A078922	001	Feb 14, 2008

PRESCRIPTION DRUG PRODUCT LIST

ZILEUTON

TABLET; ORAL

ZYFLO

+! CHIESI USA INC 600MG

N020471 003 Dec 09, 1996

TABLET, EXTENDED RELEASE; ORAL

ZILEUTON

AB	LUPIN LTD	600MG	A211972 001	Nov 05, 2019
AB	PAR PHARM INC	600MG	A212670 001	Dec 16, 2019
AB	RISING	600MG	A204929 001	Mar 17, 2017

ZYFLO CR

AB	+! CHIESI USA INC	600MG	N022052 001	May 30, 2007
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ZINC ACETATE

CAPSULE; ORAL

GALZIN

+ TEVA EQ 25MG ZINC

N020458 001 Jan 28, 1997

+! EQ 50MG ZINC

N020458 002 Jan 28, 1997

ZINC CHLORIDE

INJECTABLE; INJECTION

ZINC CHLORIDE IN PLASTIC CONTAINER

+! HOSPIRA EQ 1MG ZINC/ML

N018959 001 Jun 26, 1986

ZINC SULFATE

SOLUTION; INTRAVENOUS

ZINC SULFATE

+! AM REGENT EQ 25MG BASE/5ML (EQ 5MG BASE/ML)

N209377 002 Jul 18, 2019

+! EQ 30MG BASE/10ML (EQ 3MG BASE/ML)

N209377 001 Jul 18, 2019

ZIPRASIDONE HYDROCHLORIDE

CAPSULE; ORAL

GEODON

AB	+! PFIZER	EQ 20MG BASE	N020825 001	Feb 05, 2001
AB	+	EQ 40MG BASE	N020825 002	Feb 05, 2001
AB	+	EQ 60MG BASE	N020825 003	Feb 05, 2001
AB	+	EQ 80MG BASE	N020825 004	Feb 05, 2001

ZIPRASIDONE HYDROCHLORIDE

AB	APOTEX	EQ 20MG BASE	A077561 001	Mar 02, 2012
AB		EQ 40MG BASE	A077561 002	Mar 02, 2012
AB		EQ 60MG BASE	A077561 003	Mar 02, 2012
AB		EQ 80MG BASE	A077561 004	Mar 02, 2012
AB	AUROBINDO PHARMA LTD	EQ 20MG BASE	A204117 001	Dec 27, 2016
AB		EQ 40MG BASE	A204117 002	Dec 27, 2016
AB		EQ 60MG BASE	A204117 003	Dec 27, 2016
AB		EQ 80MG BASE	A204117 004	Dec 27, 2016
AB	DR REDDYS LABS INC	EQ 20MG BASE	A077565 001	Mar 02, 2012
AB		EQ 40MG BASE	A077565 002	Mar 02, 2012
AB		EQ 60MG BASE	A077565 003	Mar 02, 2012
AB		EQ 80MG BASE	A077565 004	Mar 02, 2012
AB	LUPIN PHARMS	EQ 20MG BASE	A077560 001	Mar 02, 2012
AB		EQ 40MG BASE	A077560 002	Mar 02, 2012
AB		EQ 60MG BASE	A077560 003	Mar 02, 2012
AB		EQ 80MG BASE	A077560 004	Mar 02, 2012
AB	MACLEODS PHARMS LTD	EQ 20MG BASE	A204375 001	Feb 17, 2017
AB		EQ 40MG BASE	A204375 002	Feb 17, 2017
AB		EQ 60MG BASE	A204375 003	Feb 17, 2017
AB		EQ 80MG BASE	A204375 004	Feb 17, 2017
AB	SANDOZ INC	EQ 20MG BASE	A077562 001	Jun 01, 2012
AB		EQ 40MG BASE	A077562 002	Jun 01, 2012
AB		EQ 60MG BASE	A077562 003	Jun 01, 2012
AB		EQ 80MG BASE	A077562 004	Jun 01, 2012
AB	WOCKHARDT LTD	EQ 20MG BASE	A090348 001	Sep 05, 2012
AB		EQ 40MG BASE	A090348 002	Sep 05, 2012
AB		EQ 60MG BASE	A090348 003	Sep 05, 2012
AB		EQ 80MG BASE	A090348 004	Sep 05, 2012

ZIPRASIDONE MESYLATE

INJECTABLE; INTRAMUSCULAR

GEODON

AP	+! PFIZER	EQ 20MG BASE/ML	N020919 001	Jun 21, 2002
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ZIPRASIDONE MESYLATE

AP	GLAND PHARMA LTD	EQ 20MG BASE/ML	A211908 001	Dec 26, 2019
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PRESCRIPTION DRUG PRODUCT LIST

3-451 (of 453)

ZOLEDRONIC ACID

INJECTABLE; INTRAVENOUS

RECLAST

<u>AP</u>	<u>+!</u> NOVARTIS	<u>EQ 5MG BASE/100ML</u>	<u>N021817 001</u>	Apr 16, 2007
<u>ZOLEDRONIC</u>				
<u>AP</u>	GLAND PHARMA LTD	<u>EQ 4MG BASE/100ML</u>	<u>A205749 001</u>	Jun 29, 2018
<u>ZOLEDRONIC ACID</u>				
<u>AP</u>	ACCORD HLTHCARE	<u>EQ 4MG BASE/5ML</u>	<u>A205279 001</u>	Nov 28, 2016
<u>AP</u>	AKORN	<u>EQ 5MG BASE/100ML</u>	<u>A200918 001</u>	Aug 21, 2014
<u>AP</u>	AKORN INC	<u>EQ 4MG BASE/5ML</u>	<u>A202548 001</u>	May 22, 2014
<u>AP</u>	APOTEX	<u>EQ 5MG BASE/100ML</u>	<u>A204367 001</u>	Dec 24, 2015
<u>AP</u>	AUROBINDO PHARMA LTD	<u>EQ 4MG BASE/5ML</u>	<u>A207751 001</u>	Sep 26, 2016
<u>AP</u>		<u>EQ 5MG BASE/100ML</u>	<u>A209125 001</u>	Dec 08, 2017
<u>AP</u>	BPI LABS LLC	<u>EQ 4MG BASE/5ML</u>	<u>A207341 001</u>	Dec 29, 2017
<u>AP</u>	BRECKENRIDGE	<u>EQ 4MG BASE/5ML</u>	<u>A091170 001</u>	Mar 04, 2013
<u>AP</u>		<u>EQ 4MG BASE/5ML</u>	<u>A202571 001</u>	May 07, 2013
<u>AP</u>		<u>EQ 5MG BASE/100ML</u>	<u>A202163 001</u>	Aug 05, 2013
<u>AP</u>	CIPLA	<u>EQ 4MG BASE/100ML</u>	<u>A210174 001</u>	Oct 27, 2017
<u>AP</u>	DR REDDYS LABS LTD	<u>EQ 4MG BASE/5ML</u>	<u>A091186 001</u>	Mar 04, 2013
<u>AP</u>		<u>EQ 5MG BASE/100ML</u>	<u>A091363 001</u>	Mar 29, 2013
<u>AP</u>	EMCURE PHARMS LTD	<u>EQ 4MG BASE/5ML</u>	<u>A201783 001</u>	Mar 12, 2013
<u>AP</u>		<u>EQ 5MG BASE/100ML</u>	<u>A201801 001</u>	Mar 29, 2013
<u>AP</u>	FRESENIUS KABI USA	<u>EQ 4MG BASE/5ML</u>	<u>A091516 001</u>	Apr 23, 2015
<u>AP</u>	GLAND PHARMA LTD	<u>EQ 4MG BASE/5ML</u>	<u>A202930 001</u>	Aug 05, 2013
<u>AP</u>		<u>EQ 5MG BASE/100ML</u>	<u>A204217 001</u>	Aug 18, 2016
<u>AP</u>		<u>EQ 5MG BASE/100ML</u>	<u>A209578 001</u>	Aug 08, 2019
<u>AP</u>	HIKMA FARMACEUTICA	<u>EQ 4MG BASE/5ML</u>	<u>A202182 001</u>	Jun 03, 2013
<u>AP</u>	HOSPIRA INC	<u>EQ 4MG BASE/5ML</u>	<u>A090621 001</u>	Mar 19, 2015
<u>AP</u>		<u>EQ 5MG BASE/100ML</u>	<u>A202837 001</u>	Apr 05, 2013
<u>AP</u>	INFORLIFE	<u>EQ 4MG BASE/100ML</u>	<u>N203231 001</u>	Aug 02, 2013
<u>AP</u>		<u>EQ 5MG BASE/100ML</u>	<u>A202828 001</u>	Sep 23, 2013
<u>AP</u>	INGENUS PHARMS LLC	<u>EQ 4MG BASE/5ML</u>	<u>A208968 001</u>	Feb 19, 2020
<u>AP</u>	MYLAN LABS LTD	<u>EQ 4MG BASE/5ML</u>	<u>A202650 001</u>	Mar 04, 2013
<u>AP</u>		<u>EQ 5MG BASE/100ML</u>	<u>A203841 001</u>	Feb 14, 2017
<u>AP</u>		<u>EQ 5MG BASE/100ML</u>	<u>A205254 001</u>	Oct 27, 2017
<u>AP</u>	SAGENT PHARMS INC	<u>EQ 4MG BASE/5ML</u>	<u>A091493 001</u>	Nov 24, 2014
<u>AP</u>	USV	<u>EQ 4MG BASE/5ML</u>	<u>A202923 001</u>	Sep 04, 2014

ZOMETA

<u>AP</u>	<u>+!</u> NOVARTIS	<u>EQ 4MG BASE/5ML</u>	<u>N021223 002</u>	Mar 07, 2003
<u>AP</u>	<u>+!</u>	<u>EQ 4MG BASE/100ML</u>	<u>N021223 003</u>	Jun 17, 2011

SOLUTION; INTRAVENOUS

ZOLEDRONIC ACID

HOSPIRA INC	EQ 4MG BASE/100ML (EQ 0.04MG BASE/ML)	N204016 001	Dec 28, 2015
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ZOLMITRIPTAN

SPRAY; NASAL

ZOMIG

<u>+</u> ASTRAZENECA	2.5MG/SPRAY	N021450 003	Sep 16, 2013
<u>+!</u>	5MG/SPRAY	N021450 004	Sep 30, 2003

TABLET; ORAL

ZOLMITRIPTAN

<u>AB</u>	AJANTA PHARMA LTD	<u>2.5MG</u>	<u>A204041 001</u>	May 20, 2016
<u>AB</u>		<u>5MG</u>	<u>A204041 002</u>	May 20, 2016
<u>AB</u>	ALEMBIC PHARMS LTD	<u>2.5MG</u>	<u>A204232 001</u>	Sep 30, 2015
<u>AB</u>		<u>5MG</u>	<u>A204232 002</u>	Sep 30, 2015
<u>AB</u>	AUROBINDO PHARMA LTD	<u>2.5MG</u>	<u>A207021 001</u>	May 11, 2016
<u>AB</u>		<u>5MG</u>	<u>A207021 002</u>	May 11, 2016
<u>AB</u>	GLENMARK GENERICS	<u>2.5MG</u>	<u>A201779 001</u>	May 14, 2013
<u>AB</u>		<u>5MG</u>	<u>A201779 002</u>	May 14, 2013
<u>AB</u>	INVAGEN PHARMS	<u>2.5MG</u>	<u>A204284 001</u>	Apr 09, 2014
<u>AB</u>		<u>5MG</u>	<u>A204284 002</u>	Apr 09, 2014
<u>AB</u>	JUBILANT GENERICS	<u>2.5MG</u>	<u>A202279 001</u>	Nov 20, 2014
<u>AB</u>		<u>5MG</u>	<u>A202279 002</u>	Nov 20, 2014
<u>AB</u>	ORCHID HLTHCARE	<u>2.5MG</u>	<u>A203726 001</u>	Oct 21, 2019
<u>AB</u>		<u>5MG</u>	<u>A203726 002</u>	Oct 21, 2019
<u>AB</u>	PLD ACQUISITIONS LLC	<u>2.5MG</u>	<u>A207867 001</u>	Feb 27, 2017
<u>AB</u>		<u>5MG</u>	<u>A207867 002</u>	Feb 27, 2017
<u>AB</u>	TWI PHARMS	<u>2.5MG</u>	<u>A206973 001</u>	Jun 30, 2017
<u>AB</u>		<u>5MG</u>	<u>A206973 002</u>	Jun 30, 2017
<u>AB</u>	ZYDUS PHARMS	<u>2.5MG</u>	<u>A203019 001</u>	Jul 11, 2018

PRESCRIPTION DRUG PRODUCT LIST

ZOLMITRIPTAN

TABLET;ORAL

ZOLMITRIPTAN

<u>AB</u>		<u>5MG</u>	<u>A203019</u>	<u>002</u>	Jul 11, 2018
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ZOMIG

<u>AB</u>	+	IPR	<u>2.5MG</u>	<u>N020768</u>	<u>001</u>	Nov 25, 1997
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<u>AB</u>	+	!	<u>5MG</u>	<u>N020768</u>	<u>002</u>	Nov 25, 1997
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TABLET, ORALLY DISINTEGRATING;ORAL

ZOLMITRIPTAN

<u>AB</u>		ALEMBIC PHARMS LTD	<u>2.5MG</u>	<u>A205074</u>	<u>001</u>	Dec 01, 2016
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<u>AB</u>			<u>5MG</u>	<u>A205074</u>	<u>002</u>	Dec 01, 2016
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<u>AB</u>		GLENMARK GENERICS	<u>2.5MG</u>	<u>A202560</u>	<u>001</u>	May 14, 2013
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<u>AB</u>			<u>5MG</u>	<u>A202560</u>	<u>002</u>	May 14, 2013
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<u>AB</u>		JUBILANT GENERICS	<u>2.5MG</u>	<u>A202956</u>	<u>001</u>	Sep 17, 2015
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<u>AB</u>			<u>5MG</u>	<u>A202956</u>	<u>002</u>	Sep 17, 2015
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<u>AB</u>		ZYDUS PHARMS USA	<u>2.5MG</u>	<u>A202890</u>	<u>001</u>	May 15, 2013
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<u>AB</u>		INC	<u>5MG</u>	<u>A202890</u>	<u>002</u>	May 15, 2013
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ZOMIG-ZMT

<u>AB</u>	+	ASTRAZENECA	<u>2.5MG</u>	<u>N021231</u>	<u>001</u>	Feb 13, 2001
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<u>AB</u>	+	!	<u>5MG</u>	<u>N021231</u>	<u>002</u>	Sep 17, 2001
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ZOLPIDEM TARTRATE

SPRAY, METERED;ORAL

ZOLPIMIST

+	!	AYTU	5MG/SPRAY	N022196	001	Dec 19, 2008
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TABLET;ORAL

AMBIEN

<u>AB</u>	+	SANOFI AVENTIS US	<u>5MG</u>	<u>N019908</u>	<u>001</u>	Dec 16, 1992
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<u>AB</u>	+	!	<u>10MG</u>	<u>N019908</u>	<u>002</u>	Dec 16, 1992
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ZOLPIDEM TARTRATE

<u>AB</u>		ACME LABS	<u>5MG</u>	<u>A077214</u>	<u>001</u>	Apr 23, 2007
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<u>AB</u>			<u>10MG</u>	<u>A077214</u>	<u>002</u>	Apr 23, 2007
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<u>AB</u>		APOTEX INC	<u>5MG</u>	<u>A077884</u>	<u>001</u>	Apr 23, 2007
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<u>AB</u>			<u>10MG</u>	<u>A077884</u>	<u>002</u>	Apr 23, 2007
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<u>AB</u>		AUROBINDO PHARMA	<u>5MG</u>	<u>A078413</u>	<u>001</u>	May 04, 2007
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<u>AB</u>			<u>10MG</u>	<u>A078413</u>	<u>002</u>	May 04, 2007
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<u>AB</u>		CIPLA LTD	<u>5MG</u>	<u>A077388</u>	<u>001</u>	Jul 30, 2012
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<u>AB</u>			<u>10MG</u>	<u>A077388</u>	<u>002</u>	Jul 30, 2012
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<u>AB</u>		INVAGEN PHARMS	<u>5MG</u>	<u>A078184</u>	<u>001</u>	Sep 07, 2007
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<u>AB</u>			<u>10MG</u>	<u>A078184</u>	<u>002</u>	Sep 07, 2007
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<u>AB</u>		SANDOZ INC	<u>5MG</u>	<u>A077322</u>	<u>001</u>	Apr 23, 2007
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<u>AB</u>			<u>10MG</u>	<u>A077322</u>	<u>002</u>	Apr 23, 2007
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<u>AB</u>		TEVA	<u>5MG</u>	<u>A076410</u>	<u>001</u>	Apr 23, 2007
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<u>AB</u>			<u>10MG</u>	<u>A076410</u>	<u>002</u>	Apr 23, 2007
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<u>AB</u>		TORRENT PHARMS	<u>5MG</u>	<u>A077903</u>	<u>001</u>	Aug 17, 2007
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<u>AB</u>			<u>10MG</u>	<u>A077903</u>	<u>002</u>	Aug 17, 2007
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<u>AB</u>		VINTAGE	<u>5MG</u>	<u>A078616</u>	<u>001</u>	Nov 21, 2008
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<u>AB</u>			<u>10MG</u>	<u>A078616</u>	<u>002</u>	Nov 21, 2008
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<u>AB</u>		YUNG SHIN PHARM	<u>5MG</u>	<u>A077990</u>	<u>001</u>	Apr 23, 2007
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<u>AB</u>			<u>10MG</u>	<u>A077990</u>	<u>002</u>	Apr 23, 2007
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TABLET;SUBLINGUAL

EDLUAR

<u>AB</u>	+	MYLAN SPECIALITY LP	<u>5MG</u>	<u>N021997</u>	<u>001</u>	Mar 13, 2009
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<u>AB</u>	+	!	<u>10MG</u>	<u>N021997</u>	<u>002</u>	Mar 13, 2009
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ZOLPIDEM TARTRATE

<u>AB</u>		DR REDDYS	<u>1.75MG</u>	<u>A204503</u>	<u>001</u>	Nov 18, 2016
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<u>AB</u>			<u>3.5MG</u>	<u>A204503</u>	<u>002</u>	Nov 18, 2016
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<u>AB</u>		MYLAN	<u>5MG</u>	<u>A202657</u>	<u>001</u>	Aug 08, 2016
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<u>AB</u>			<u>10MG</u>	<u>A202657</u>	<u>002</u>	Aug 08, 2016
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<u>AB</u>		NOVEL LABS INC	<u>1.75MG</u>	<u>A204299</u>	<u>001</u>	Jun 03, 2015
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<u>AB</u>	!		<u>3.5MG</u>	<u>A204299</u>	<u>002</u>	Jun 03, 2015
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<u>AB</u>		PAR FORM	<u>5MG</u>	<u>A201509</u>	<u>001</u>	Aug 01, 2016
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<u>AB</u>			<u>10MG</u>	<u>A201509</u>	<u>002</u>	Aug 01, 2016
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<u>AB</u>		PAR PHARM INC	<u>1.75MG</u>	<u>A204229</u>	<u>001</u>	Sep 11, 2017
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<u>AB</u>			<u>3.5MG</u>	<u>A204229</u>	<u>002</u>	Sep 11, 2017
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TABLET, EXTENDED RELEASE;ORAL

AMBIEN CR

<u>AB</u>	+	SANOFI AVENTIS US	<u>6.25MG</u>	<u>N021774</u>	<u>002</u>	Sep 02, 2005
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<u>AB</u>	+	!	<u>12.5MG</u>	<u>N021774</u>	<u>001</u>	Sep 02, 2005
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ZOLPIDEM TARTRATE

<u>AB</u>		ACTAVIS ELIZABETH	<u>6.25MG</u>	<u>A078179</u>	<u>002</u>	Oct 13, 2010
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<u>AB</u>			<u>12.5MG</u>	<u>A078179</u>	<u>001</u>	Jun 06, 2011
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PRESCRIPTION DRUG PRODUCT LIST

ZOLPIDEM TARTRATE

TABLET, EXTENDED RELEASE;ORAL

ZOLPIDEM TARTRATE

<u>AB</u>	ACTAVIS LABS FL INC	<u>6.25MG</u>	<u>A090153 001</u>	Mar 25, 2013
<u>AB</u>		<u>12.5MG</u>	<u>A090153 002</u>	Mar 25, 2013
<u>AB</u>	ANCHEN PHARMS	<u>6.25MG</u>	<u>A078148 002</u>	Apr 14, 2011
<u>AB</u>		<u>12.5MG</u>	<u>A078148 001</u>	Dec 03, 2010
<u>AB</u>	APOTEX INC	<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>	+ SUNOVION PHARMS INC	<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>	BIONPHARMA INC	<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>	CELLTRION	<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>	GLENMARK GENERICS	<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>	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>	ZYDUS PHARMS USA	<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

OTC DRUG PRODUCT LISTACETAMINOPHEN

SUPPOSITORY;RECTAL

ACEPHEN

ACP NIMBLE	120MG	N018060 001	
	325MG	A072344 001	Mar 27, 1992
	650MG	A072237 001	Mar 27, 1992

ACETAMINOPHEN

PERRIGO NEW YORK	120MG	A070607 001	Apr 06, 1987
	650MG	A070608 001	Dec 01, 1986
+ TARO PHARM INDS LTD	120MG	N018337 003	Sep 12, 1983
+	325MG	N018337 002	
+!	650MG	N018337 001	

INFANTS' FEVERALL

+ TARO PHARM INDS LTD	80MG	N018337 004	Aug 26, 1992
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NEOPAP

POLYMEDICA	120MG	N016401 001	
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TABLET, EXTENDED RELEASE;ORAL

ACETAMINOPHEN

AUROBINDO PHARMA LTD	650MG	A207229 001	Nov 09, 2016
GRANULES INDIA LTD	650MG	A211544 001	Apr 16, 2019
HERITAGE PHARMA	650MG	A207035 001	May 31, 2018
OHM LABS	650MG	A076200 001	Mar 19, 2002
PERRIGO	650MG	A075077 001	Feb 25, 2000
SUN PHARM INDS LTD	650MG	A078569 001	Dec 14, 2011

TYLENOL

+! J AND J CONSUMER INC	650MG	N019872 001	Jun 08, 1994
+!	650MG	N019872 002	Jan 11, 2001

ACETAMINOPHEN; ASPIRIN; CAFFEINE

TABLET;ORAL

ACETAMINOPHEN, ASPIRIN AND CAFFEINE

PERRIGO	250MG;250MG;65MG	A075794 001	Nov 26, 2001
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EXCEDRIN (MIGRAINE)

+! GLAXOSMITHKLINE CONS	250MG;250MG;65MG	N020802 001	Jan 14, 1998
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ACETAMINOPHEN; IBUPROFEN

TABLET;ORAL

ADVIL DUAL ACTION WITH ACETAMINOPHEN

+! PFIZER INC	250MG;125MG	N211733 001	Feb 28, 2020
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ADAPALENE

GEL;TOPICAL

ADAPALENE

GLENMARK GENERICS	0.1%	A091314 002	Jul 09, 2019
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DIFFERIN

+! GALDERMA LABS LP	0.1%	N020380 002	Jul 08, 2016
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ALCOHOL; CHLORHEXIDINE GLUCONATE

SOLUTION;TOPICAL

AVAGARD

+! 3M	61%;1%	N021074 001	Jun 07, 2001
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ALUMINUM HYDROXIDE; MAGNESIUM TRISILICATE

TABLET, CHEWABLE;ORAL

GAVISCON

+ SANOFI AVENTIS US	80MG;20MG	N018685 001	Dec 09, 1983
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ASPIRIN

CAPSULE;ORAL

VAZALORE

+! PLX PHARMA	325MG	N203697 001	Jan 14, 2013
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AVOBENZONE; ECAMSULE; OCTOCRYLENE

CREAM;TOPICAL

ANTHELIOS SX

+! LOREAL USA	2%;2%;10%	N021502 001	Jul 21, 2006
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CAPITAL SOLEIL 15

+! LOREAL USA	2%;3%;10%	N021501 001	Oct 02, 2006
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OTC DRUG PRODUCT LISTAVOBENZONE; 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

BENTOQUATAM

LOTION; TOPICAL

IVY BLOCK

+! STAND HOMEOPATH 5% N020532 001 Aug 26, 1996

BRIMONIDINE TARTRATE

SOLUTION/DROPS; OPHTHALMIC

LUMIFY

+! BAUSCH AND LOMB INC 0.025% N208144 001 Dec 22, 2017

BUDESONIDE

SPRAY, METERED; NASAL

BUDESONIDE

APOTEX INC 0.032MG/SPRAY A078949 002 Nov 20, 2015

RHINOCORT ALLERGY

+! ASTRAZENECA PHARMS 0.032MG/SPRAY N020746 003 Mar 23, 2015

BUTENAFINE HYDROCHLORIDE

CREAM; TOPICAL

BUTENAFINE HYDROCHLORIDE

TARO PHARMS 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

800MG;10MG;165MG A204782 001 Aug 29, 2016

PEPCID COMPLETE

+! J AND J CONSUMER 800MG;10MG;165MG N020958 001 Oct 16, 2000

INC

CETIRIZINE HYDROCHLORIDE

CAPSULE; ORAL

CETIRIZINE HYDROCHLORIDE ALLERGY

APOTEX 10MG A207235 001 Aug 12, 2016

AUROBINDO PHARMA 10MG A209107 001 Jul 20, 2018

LTD

+ BIONPHARMA INC 5MG N022429 001 Jul 23, 2009

+! 10MG N022429 004 Jul 23, 2009

STRIDES PHARMA 10MG A205291 001 Jul 21, 2017

CETIRIZINE HYDROCHLORIDE HIVES RELIEF

AUROBINDO PHARMA 10MG A209107 002 Jul 20, 2018

LTD

+ BIONPHARMA INC 5MG N022429 003 Jul 23, 2009

+! 10MG N022429 002 Jul 23, 2009

SYRUP; ORAL

CHILDREN'S CETIRIZINE HYDROCHLORIDE ALLERGY

AMNEAL PHARMS 5MG/5ML A090765 002 Oct 07, 2009

ATHEM 5MG/5ML A091327 001 Oct 17, 2011

AUROBINDO PHARMA 5MG/5ML A090750 002 Feb 02, 2010

HETERO LABS LTD III 5MG/5ML A210622 001 Mar 13, 2019

LANNETT CO INC 5MG/5ML A091130 001 Apr 22, 2011

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

TRIS PHARMA INC 5MG/5ML A090572 001 Nov 16, 2012

CHILDREN'S CETIRIZINE HYDROCHLORIDE HIVES RELIEF

AMNEAL PHARMS 5MG/5ML A090765 001 Oct 07, 2009

ATHEM 5MG/5ML A091327 002 Oct 17, 2011

AUROBINDO PHARMA 5MG/5ML A090750 001 Feb 02, 2010

LANNETT CO INC 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

OTC DRUG PRODUCT LIST

CETIRIZINE HYDROCHLORIDE

SYRUP; ORAL

CHILDREN'S CETIRIZINE HYDROCHLORIDE HIVES RELIEF

	5MG/5ML	A201546	002	May 20, 2011
TRIS PHARMA INC	5MG/5ML	A090572	002	Nov 16, 2012
CHILDREN'S ZYRTEC ALLERGY				
+! J AND J CONSUMER	5MG/5ML	N022155	002	Nov 16, 2007
INC				
CHILDREN'S ZYRTEC HIVES RELIEF				
+! J AND J CONSUMER	5MG/5ML	N022155	001	Nov 16, 2007
INC				

TABLET; ORAL

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 LTD	5MG	A090760	001	Aug 05, 2015
	10MG	A090760	003	Aug 05, 2015
CIPLA LTD	5MG	A077318	001	Jul 25, 2013
	10MG	A077318	002	Jul 25, 2013
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
GRANULES INDIA LTD	10MG	A209274	001	Dec 22, 2017
IPCA LABS LTD	5MG	A202277	002	Mar 11, 2014
	10MG	A202277	004	Mar 11, 2014
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
ORCHID HLTHCARE	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
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
UNIQUE PHARM LABS	5MG	A077829	001	Aug 26, 2009
	10MG	A077829	004	Aug 26, 2009
WOCKHARDT	5MG	A078427	003	Dec 28, 2007
	10MG	A078427	004	Dec 28, 2007
CETIRIZINE HYDROCHLORIDE HIVES				
DR REDDYS LABS LTD	5MG	A078343	001	Jan 15, 2008
	10MG	A078343	002	Jan 15, 2008
IPCA LABS LTD	5MG	A202277	001	Mar 11, 2014
	10MG	A202277	003	Mar 11, 2014
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
ORCHID HLTHCARE	5MG	A078862	003	Feb 19, 2009
	10MG	A078862	004	Feb 19, 2009
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
UNICHEM	5MG	A078680	001	Jun 26, 2009
	10MG	A078680	002	Jun 26, 2009
UNIQUE PHARM LABS	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

OTC DRUG PRODUCT LISTCETIRIZINE HYDROCHLORIDE

TABLET;ORAL

CETIRIZINE HYDROCHLORIDE HIVES RELIEF

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
TORRENT PHARMS LLC	5MG	A079191 003	Apr 15, 2010
	10MG	A079191 002	Apr 15, 2010

ZYRTEC ALLERGY

+! J AND J CONSUMER INC	10MG	N019835 004	Nov 16, 2007
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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
SUN PHARM	5MG	A090142 001	Aug 30, 2011
	10MG	A090142 002	Aug 30, 2011

CHILDREN'S CETIRIZINE HYDROCHLORIDE HIVES RELIEF

JUBILANT GENERICS	5MG	A091116 003	Feb 19, 2015
	10MG	A091116 004	Feb 19, 2015
SUN PHARM	5MG	A090142 003	Aug 30, 2011
	10MG	A090142 004	Aug 30, 2011

TABLET, ORALLY DISINTEGRATING;ORAL

CETIRIZINE HYDROCHLORIDE ALLERGY

PERRIGO R AND D	10MG	A205490 001	Sep 02, 2015
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ZYRTEC ALLERGY

+! J AND J CONSUMER INC	10MG	N022578 001	Sep 03, 2010
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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
SUN PHARM INDS LTD	5MG;120MG	A090922 001	Sep 28, 2012

ZYRTEC-D 12 HOUR

+! J AND J CONSUMER INC	5MG;120MG	N021150 002	Nov 09, 2007
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CHLORHEXIDINE GLUCONATE

AEROSOL, METERED;TOPICAL

EXIDINE

+! XTTRIUM	4%	N019127 001	Dec 24, 1984
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CLOTH;TOPICAL

CHLORHEXIDINE GLUCONATE

+! SAGE PRODS	2%	N021669 001	Apr 25, 2005
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READYPREP CHG

MEDLINE INDUSTRIES	2%	N207964 001	Nov 20, 2018
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SOLUTION;TOPICAL

BRIAN CARE

SOAPCO	4%	A071419 001	Dec 17, 1987
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CHG SCRUB

ECOLAB	4%	N019258 002	Jul 22, 1986
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CHLORHEXIDINE GLUCONATE

BAJAJ	0.75%	N020111 001	Sep 11, 1997
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CIDA-STAT

ECOLAB	2%	N019258 001	Jul 22, 1986
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EXIDINE

+! XTTRIUM	2%	N019422 001	Dec 17, 1985
	4%	N019125 001	Dec 24, 1984

HIBICLENS

+! MOLNLYCKE HLTH	4%	N017768 001	
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HIBISTAT

+! MOLNLYCKE HLTH	0.5%	N018300 001	
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SPONGE;TOPICAL

BIOSCRUB

GRIFFEN	4%	N019822 001	Mar 31, 1989
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OTC DRUG PRODUCT LIST

CHLORHEXIDINE GLUCONATE

SPONGE; TOPICAL

CHLORHEXIDINE GLUCONATE

! BECTON DICKINSON 4% A072525 001 Oct 24, 1989

CHLORHEXIDINE GLUCONATE; ISOPROPYL ALCOHOL

SOLUTION; TOPICAL

SOLUPREP

+! 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

+! GLAXOSMITHKLINE 4MG;200MG;10MG N022113 001 Dec 21, 2011

ADVIL MULTI-SYMPTOM COLD & FLU

+! GLAXOSMITHKLINE 4MG;200MG;10MG N022113 002 Apr 28, 2017

CHLORPHENIRAMINE MALEATE; IBUPROFEN; PSEUDOEPHEDRINE HYDROCHLORIDE

SUSPENSION; ORAL

CHILDREN'S ADVIL ALLERGY SINUS

+! GLAXOSMITHKLINE 1MG/5ML;100MG/5ML;15MG/5ML N021587 001 Feb 24, 2004

TABLET; ORAL

ADVIL ALLERGY SINUS

+! GLAXOSMITHKLINE 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

IVAX SUB TEVA 200MG A075345 001 Jun 16, 1999

PHARMS L PERRIGO CO 200MG A075285 001 Oct 29, 1998

TAGAMET HB

+! MEDTECH PRODUCTS 200MG N020238 002 Aug 21, 1996

CLEMASTINE FUMARATE

TABLET; ORAL

CLEMASTINE FUMARATE

! L PERRIGO CO 1.34MG A074512 001 Nov 22, 1995

OTC DRUG PRODUCT LISTCLOTRIMAZOLE

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
	PERRIGO	5.2MG/SPRAY	A075427	001	Dec 12, 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 LTD	30MG; 600MG	A206941	001	Mar 17, 2017
		60MG; 1.2GM	A206941	002	Mar 17, 2017
	PERRIGO R AND D	30MG; 600MG	A207602	002	Mar 05, 2018
		60MG; 1.2GM	A207602	001	Mar 05, 2018
	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
		AMNEAL PHARMS LLC	EQ 30MG HYDROBROMIDE/5ML	A203133	001	Jul 28, 2017
		TRIS PHARMA INC	EQ 30MG HYDROBROMIDE/5ML	A091135	001	May 25, 2012

DIPHENHYDRAMINE CITRATE; IBUPROFEN

TABLET; ORAL

ADVIL PM

+	!	GLAXOSMITHKLINE	38MG; 200MG	N021394	001	Dec 21, 2005
		IBUPROFEN AND DIPHENHYDRAMINE CITRATE				
		DR REDDYS LABS LTD	38MG; 200MG	A090619	001	Jul 08, 2009
		PERRIGO R AND D	38MG; 200MG	A079113	001	Dec 22, 2008

DIPHENHYDRAMINE HYDROCHLORIDE; IBUPROFEN

CAPSULE; ORAL

ADVIL PM

+	!	GLAXOSMITHKLINE	25MG; EQ 200MG FREE ACID AND POTASSIUM SALT	N021393	001	Dec 21, 2005
		IBUPROFEN AND DIPHENHYDRAMINE HYDROCHLORIDE				
		AUROBINDO PHARMA LTD	25MG; EQ 200MG FREE ACID AND POTASSIUM SALT	A210676	001	Feb 14, 2019
		BIONPHARMA INC	25MG; EQ 200MG FREE ACID AND POTASSIUM SALT	A090397	001	Nov 22, 2010
		STRIDES PHARMA	25MG; EQ 200MG FREE ACID AND POTASSIUM SALT	A200888	001	Mar 05, 2012

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
		APOTEX	25MG; 220MG	A211830	001	Aug 22, 2019
		PERRIGO R AND D	25MG; 220MG	A208499	001	May 10, 2019

DOCOSANOL

CREAM; TOPICAL

ABREVA

+	!	GLAXOSMITHKLINE	10%	N020941	001	Jul 25, 2000
		DOCOSANOL				
		P AND L	10%	A208754	001	Nov 19, 2018

OTC DRUG PRODUCT LIST

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	
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EPINEPHRINE

AEROSOL, METERED; INHALATION

PRIMATENE MIST

+! ARMSTRONG PHARMS	0.125MG/INH	N205920	001	Nov 07, 2018
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ESOMEPRAZOLE MAGNESIUM

CAPSULE, DELAYED RELEASE;ORAL

ESOMEPRAZOLE MAGNESIUM

AMNEAL PHARMS NY	EQ 20MG BASE	A209716	001	Jun 05, 2019
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AUROBINDO PHARMA LTD	EQ 20MG BASE	A209339	001	Oct 16, 2017
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DR REDDYS LABS LTD	EQ 20MG BASE	A207673	001	May 15, 2018
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MYLAN	EQ 20MG BASE	A212376	001	Oct 16, 2019
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PERRIGO R AND D	EQ 20MG BASE	A207193	001	Aug 18, 2017
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NEXIUM 24HR

+! ASTRAZENECA LP	EQ 20MG BASE	N204655	001	Mar 28, 2014
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TABLET, DELAYED RELEASE;ORAL

NEXIUM 24HR

+! ASTRAZENECA LP	EQ 20MG BASE	N207920	001	Nov 23, 2015
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FAMOTIDINE

TABLET;ORAL

FAMOTIDINE

AUROBINDO PHARMA LTD	10MG	A206531	001	Apr 26, 2016
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	20MG	A206531	002	Apr 26, 2016
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DR REDDYS LABS LTD	10MG	A077367	002	Aug 17, 2001
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	20MG	A077367	001	Sep 25, 2006
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P AND L	10MG	A075512	001	Jul 26, 2001
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PERRIGO	10MG	A075400	001	Mar 18, 2005
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PERRIGO R AND D	20MG	A077351	001	Sep 25, 2006
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SUN PHARM INDS LTD	10MG	A090283	001	Nov 17, 2009
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	20MG	A090283	002	Nov 17, 2009
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TEVA	10MG	A075312	001	May 31, 2001
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WOCKHARDT	10MG	A077146	001	Mar 07, 2005
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	20MG	A090837	001	Aug 04, 2010
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PEPCID AC

+ J AND J CONSUMER INC	10MG	N020325	001	Apr 28, 1995
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	10MG	N020902	001	Aug 05, 1999
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+!	20MG	N020325	002	Sep 23, 2003
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TABLET, CHEWABLE;ORAL

FAMOTIDINE

PERRIGO	10MG	A075715	001	Aug 22, 2003
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PEPCID AC

+! J AND J CONSUMER INC	20MG	N020801	002	Dec 17, 2007
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FEXOFENADINE HYDROCHLORIDE

SUSPENSION;ORAL

CHILDREN'S ALLEGRA ALLERGY

+! SANOFI AVENTIS US	30MG/5ML	N201373	001	Jan 24, 2011
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CHILDREN'S FEXOFENADINE HYDROCHLORIDE ALLERGY

P AND L	30MG/5ML	A203330	001	Nov 18, 2014
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CHILDREN'S FEXOFENADINE HYDROCHLORIDE HIVES

! P AND L	30MG/5ML	A203330	002	Nov 18, 2014
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TABLET;ORAL

ALLEGRA ALLERGY

+ SANOFI AVENTIS US	60MG	N020872	007	Jan 24, 2011
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+!	180MG	N020872	010	Jan 24, 2011
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CHILDREN'S FEXOFENADINE HYDROCHLORIDE ALLERGY

AUROLIFE PHARMA LLC	30MG	A202039	001	Nov 19, 2014
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DR REDDYS LABS LTD	30MG	A076502	004	Apr 12, 2011
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HETERO LABS LTD V	30MG	A204097	001	Aug 19, 2016
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SUN PHARM INDS	30MG	A091567	002	Feb 06, 2012
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TEVA	30MG	A076447	004	Apr 13, 2011
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WOCKHARDT LTD	30MG	A079112	002	Feb 08, 2012
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OTC DRUG PRODUCT LISTFEXOFENADINE HYDROCHLORIDE

TABLET; ORAL

CHILDREN'S FEXOFENADINE HYDROCHLORIDE HIVES

DR REDDYS LABS LTD	30MG	A076502	005	Apr 12, 2011
SUN PHARM INDS	30MG	A091567	001	Feb 06, 2012
TEVA	30MG	A076447	005	Apr 13, 2011
WOCKHARDT LTD	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

AUROLIFE PHARMA LLC	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 INDIA LTD	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
MYLAN	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	60MG	A091567	004	Feb 06, 2012
	180MG	A091567	006	Feb 06, 2012
TEVA	60MG	A076447	006	Apr 13, 2011
	180MG	A076447	008	Apr 13, 2011
UNIQUE PHARM LABS	180MG	A210137	001	Aug 13, 2018
WOCKHARDT LTD	60MG	A079112	004	Feb 08, 2012
	180MG	A079112	006	Feb 08, 2012

FEXOFENADINE HYDROCHLORIDE HIVES

DR REDDYS LABS LTD	60MG	A076502	007	Apr 12, 2011
	180MG	A076502	009	Apr 12, 2011
MYLAN	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	60MG	A091567	003	Feb 06, 2012
	180MG	A091567	005	Feb 06, 2012
TEVA	60MG	A076447	007	Apr 13, 2011
WOCKHARDT LTD	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

+! SANOFI AVENTIS US	60MG;120MG	N020786	002	Jan 24, 2011
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ALLEGRA-D 24 HOUR ALLERGY AND CONGESTION

+! SANOFI AVENTIS US	180MG;240MG	N021704	002	Jan 24, 2011
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FEXOFENADINE HYDROCHLORIDE AND PSEUDOEPHEDRINE HYDROCHLORIDE

AUROBINDO PHARMA LTD	60MG;120MG	A209116	001	Oct 30, 2017
DR REDDYS LABS LTD	60MG;120MG	A076667	001	Nov 18, 2014
	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

+! GLAXOSMITHKLINE	0.0275MG/SPRAY	N022051	002	Aug 02, 2016
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CONS

FLUTICASONE PROPIONATE

SPRAY, METERED; NASAL

FLONASE ALLERGY RELIEF

+! GLAXOSMITHKLINE	0.05MG/SPRAY	N205434	001	Jul 23, 2014
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CONS

FLUTICASONE PROPIONATE

APOTEX	0.05MG/SPRAY	A208150	001	Feb 29, 2016
HI TECH	0.05MG/SPRAY	A208024	001	Apr 17, 2019
HIKMA	0.05MG/SPRAY	A207957	001	May 26, 2016

OTC DRUG PRODUCT LIST

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 LTD	600MG	A210453 001	Oct 21, 2019
	1.2GM	A210453 002	Oct 21, 2019
GUARDIAN DRUG	600MG	A209215 001	Sep 06, 2017
	1.2GM	A209215 002	Sep 06, 2017
PERRIGO R AND D	600MG	A078912 001	Nov 23, 2011

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
DR REDDYS LABS LTD	600MG;60MG	A208369 001	Dec 29, 2017
	1.2GM;120MG	A208369 002	Dec 29, 2017

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

+! GLAXOSMITHKLINE	EQ 200MG FREE ACID AND POTASSIUM SALT	N020402 001	Apr 20, 1995
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ADVIL MIGRAINE LIQUI-GELS

+! GLAXOSMITHKLINE	EQ 200MG FREE ACID AND POTASSIUM SALT	N020402 002	Mar 16, 2000
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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
BIONPHARMA INC	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
SOFGEN PHARMS	EQ 200MG FREE ACID AND POTASSIUM SALT	A203599 001	Sep 07, 2016

MIDOL LIQUID GELS

+! BIONPHARMA INC	200MG	N021472 001	Oct 18, 2002
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SUSPENSION;ORAL

CHILDREN'S ADVIL

GLAXOSMITHKLINE	100MG/5ML	N020589 001	Jun 27, 1996
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CHILDREN'S ADVIL-FLAVORED

GLAXOSMITHKLINE	100MG/5ML	N020589 002	Nov 07, 1997
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CHILDREN'S IBUPROFEN

PERRIGO	100MG/5ML	A074937 001	Dec 22, 1998
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CHILDREN'S MOTRIN

+! J AND J CONSUMER INC	100MG/5ML	N020516 001	Jun 16, 1995
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IBUPROFEN

APTAPHARMA INC	100MG/5ML	A210602 001	Nov 23, 2018
ARISE	100MG/5ML	A200457 001	Aug 18, 2011
AUROBINDO PHARMA LTD	100MG/5ML	A209179 001	Apr 17, 2018
P AND L	100MG/5ML	A074916 001	Apr 30, 1999
TARO	100MG/5ML	A209207 001	Jun 27, 2017

SUSPENSION/DROPS;ORAL

CHILDREN'S MOTRIN

+! J AND J CONSUMER INC	40MG/ML	N020603 001	Jun 10, 1996
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IBUPROFEN

GUARDIAN DRUG	40MG/ML	A210755 001	Sep 26, 2018
L PERRIGO CO	40MG/ML	A075217 001	Dec 16, 1998
TRIS PHARMA INC	40MG/ML	A079058 001	Aug 31, 2009

INFANT'S ADVIL

+! GLAXOSMITHKLINE	50MG/1.25ML	N020812 002	Jan 12, 2000
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OTC DRUG PRODUCT LIST

4-10 (of 20)

IBUPROFEN

TABLET; ORAL

ADVIL

GLAXOSMITHKLINE 200MG N018989 001 May 18, 1984

IBU-TAB 200

ALRA 200MG A071057 001 Aug 11, 1988

IBUPROFEN

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

LTD

AVEMA PHARMA 200MG A076460 001 Nov 26, 2003

CONTRACT PHARMACAL 200MG A072299 001 Jul 01, 1988

DR REDDYS LA 200MG A075661 001 Dec 12, 2001

DR REDDYS LABS INC 200MG A076117 001 Nov 20, 2001

GRANULES INDIA 200MG A079174 001 Dec 10, 2010

GRANULES INDIA LTD 200MG A202312 001 Oct 07, 2016

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

MERRO PHARM 200MG A070985 001 Oct 02, 1987

OHM 200MG A071163 001 Jul 15, 1986

PAR PHARM 200MG A070481 001 Sep 24, 1986

PERRIGO 200MG A072096 001 Dec 08, 1987

200MG A075995 001 Mar 14, 2002

! PERRIGO R AND D 200MG A077349 001 Jun 21, 2005

SHANDONG XINHUA 200MG A206990 001 Aug 21, 2018

200MG A207095 001 May 05, 2017

STRIDES PHARMA 200MG A079129 001 Mar 28, 2011

200MG A091355 001 Apr 04, 2011

200MG A206989 001 Jun 29, 2018

200MG A207052 001 May 30, 2017

ULTRATAB LABS INC 200MG A209076 001 Jan 06, 2020

VINTAGE PHARMS 200MG A071229 001 Apr 01, 1987

200MG A071639 001 Feb 02, 1988

JUNIOR STRENGTH ADVIL

GLAXOSMITHKLINE 100MG N020267 002 Dec 13, 1996

JUNIOR STRENGTH IBUPROFEN

L PERRIGO CO 100MG A075367 001 Apr 22, 1999

JUNIOR STRENGTH MOTRIN

J AND J CONSUMER 100MG N020602 001 Jun 10, 1996

INC

MOTRIN IB

+ J AND J CONSUMER 200MG N019012 003 Dec 17, 1990

INC

TAB-PROFEN

PERRIGO 200MG A072095 001 Dec 08, 1987

TABLET, CHEWABLE; ORAL

CHILDREN'S ADVIL

GLAXOSMITHKLINE 50MG N020944 001 Dec 18, 1998

IBUPROFEN

! PERRIGO 100MG A076359 002 Jan 16, 2004

JUNIOR STRENGTH ADVIL

GLAXOSMITHKLINE 100MG N020944 002 Dec 18, 1998

IBUPROFEN SODIUM

TABLET; ORAL

ADVIL

+! GLAXOSMITHKLINE EQ 200MG BASE N201803 001 Jun 12, 2012

IBUPROFEN SODIUM

PERRIGO R AND D EQ 200MG BASE A206581 001 Aug 03, 2015

IBUPROFEN; PHENYLEPHRINE HYDROCHLORIDE

TABLET; ORAL

ADVIL CONGESTION RELIEF

+! GLAXOSMITHKLINE 200MG;10MG N022565 001 May 27, 2010

IBUPROFEN AND PHENYLEPHRINE HYDROCHLORIDE

PERRIGO R AND D 200MG;10MG A203200 001 Jul 03, 2014

OTC DRUG PRODUCT LIST

IBUPROFEN; PSEUDOEPHEDRINE HYDROCHLORIDE

CAPSULE; ORAL

ADVIL COLD AND SINUS

+	!	GLAXOSMITHKLINE	EQ 200MG FREE ACID AND POTASSIUM SALT; 30MG	N021374 001	May 30, 2002
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IBUPROFEN AND PSEUDOEPHEDRINE HYDROCHLORIDE

		AUROBINDO PHARMA LTD	EQ 200MG FREE ACID AND POTASSIUM SALT; 30MG	A209235 001	Dec 01, 2017
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SUSPENSION; ORAL

CHILDREN'S ADVIL COLD

		GLAXOSMITHKLINE	100MG/5ML; 15MG/5ML	N021373 001	Apr 18, 2002
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CHILDREN'S MOTRIN COLD

+	!	J AND J CONSUMER INC	100MG/5ML; 15MG/5ML	N021128 001	Aug 01, 2000
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IBUPROFEN AND PSEUDOEPHEDRINE HYDROCHLORIDE

		PERRIGO	100MG/5ML; 15MG/5ML	A076478 001	Nov 05, 2003
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TABLET; ORAL

ADVIL COLD AND SINUS

+	!	GLAXOSMITHKLINE	200MG; 30MG	N019771 001	Sep 19, 1989
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IBUPROFEN AND PSEUDOEPHEDRINE HYDROCHLORIDE

		DR REDDYS LABS LTD	200MG; 30MG	A077628 001	Aug 14, 2006
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IBUPROHM COLD AND SINUS

		OHM LABS	200MG; 30MG	A074567 001	Apr 17, 2001
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SINE-AID IB

		J AND J CONSUMER INC	200MG; 30MG	N019899 001	Dec 31, 1992
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INSULIN RECOMBINANT HUMAN

INJECTABLE; INJECTION

HUMULIN R

+	!	LILLY	100 UNITS/ML	N018780 001	Oct 28, 1982
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HUMULIN R PEN

+	!	LILLY	100 UNITS/ML	N018780 005	Aug 06, 1998
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NOVOLIN R

+	!	NOVO NORDISK INC	100 UNITS/ML	N019938 001	Jun 25, 1991
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INSULIN RECOMBINANT HUMAN; INSULIN SUSP ISOPHANE RECOMBINANT HUMAN

INJECTABLE; INJECTION

HUMULIN 70/30

+	!	LILLY	30 UNITS/ML; 70 UNITS/ML	N019717 001	Apr 25, 1989
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HUMULIN 70/30 PEN

+	!	LILLY	30 UNITS/ML; 70 UNITS/ML	N019717 002	Aug 06, 1998
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NOVOLIN 70/30

+	!	NOVO NORDISK INC	30 UNITS/ML; 70 UNITS/ML	N019991 001	Jun 25, 1991
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INSULIN SUSP ISOPHANE RECOMBINANT HUMAN

INJECTABLE; INJECTION

HUMULIN N

+	!	LILLY	100 UNITS/ML	N018781 001	Oct 28, 1982
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NOVOLIN N

+	!	NOVO NORDISK INC	100 UNITS/ML	N019959 001	Jul 01, 1991
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IODINE POVACRYLEX; ISOPROPYL ALCOHOL

SPONGE; TOPICAL

DURAPREP

+	!	3M	EQ 0.7% IODINE; 74% (6ML)	N021586 001	Sep 29, 2006
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+	!		EQ 0.7% IODINE; 74% (26ML)	N021586 002	Sep 29, 2006
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KETOCONAZOLE

SHAMPOO; TOPICAL

NIZORAL A-D

+	!	KRAMER	1%	N020310 001	Oct 10, 1997
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KETOTIFEN FUMARATE

SOLUTION/DROPS; OPHTHALMIC

ALAWAY

+	!	BAUSCH AND LOMB	EQ 0.025% BASE	N021996 001	Dec 01, 2006
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+			EQ 0.035% BASE	N021996 002	Feb 11, 2015
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KETOTIFEN FUMARATE

		AKORN	EQ 0.025% BASE	A077958 001	Jul 26, 2007
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		APOTEX INC	EQ 0.025% BASE	A077354 001	May 09, 2006
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ZADITOR

!		ALCON PHARMS LTD	EQ 0.025% BASE	A077200 001	Sep 02, 2008
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OTC DRUG PRODUCT LIST

LANSOPRAZOLE

CAPSULE, DELAYED REL PELLETS;ORAL

LANSOPRAZOLE

DR REDDYS LABS LTD	15MG	A202194	001	May 18, 2012
LANNETT CO INC	15MG	A207157	001	Sep 29, 2017
MYLAN	15MG	A203187	001	Jun 01, 2016
NATCO PHARMA LTD	15MG	A203306	001	Jan 13, 2016
PERRIGO R AND D	15MG	A202319	001	May 18, 2012
WOCKHARDT LTD	15MG	A202727	001	May 18, 2012

PREVACID 24 HR

+! PERRIGO PHARMA INTL	15MG	N022327	001	May 18, 2009
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TABLET, ORALLY DISINTEGRATING, DELAYED RELEASE;ORAL

LANSOPRAZOLE

DEXCEL PHARMA	15MG	N208025	001	Jun 07, 2016
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LEVOCETIRIZINE DIHYDROCHLORIDE

SOLUTION;ORAL

XYZAL ALLERGY 24HR

+! SANOFI AVENTIS US	2.5MG/5ML	N209090	001	Jan 31, 2017
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TABLET;ORAL

LEVOCETIRIZINE DIHYDROCHLORIDE

DR REDDYS LABS LTD	5MG	A210375	001	Jan 19, 2018
MICRO LABS	5MG	A211551	001	Nov 20, 2018
PERRIGO R AND D	5MG	A211983	001	Mar 28, 2019

XYZAL ALLERGY 24HR

+! SANOFI AVENTIS US	5MG	N209089	001	Jan 31, 2017
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LEVONORGESTREL

TABLET;ORAL

ATHENTIA NEXT

AUROBINDO PHARMA LTD	1.5MG	A206867	001	Dec 08, 2015
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FALLBACK SOLO

LUPIN LTD	1.5MG	A201446	001	Jun 19, 2014
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HER STYLE

NOVAST LABS	1.5MG	A207976	001	Mar 11, 2016
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LEVONORGESTREL

GLENMARK PHARMS LTD	1.5MG	A207044	001	Mar 25, 2016
! L PERRIGO CO	0.75MG	A090740	001	Dec 30, 2010
LABORATOIRE HRA	1.5MG	A204044	001	Jul 03, 2018
MYLAN LABS LTD	0.75MG	A202740	001	Sep 02, 2016
	1.5MG	A202739	001	Oct 31, 2014
NAARI PTE LTD	1.5MG	A202380	001	May 29, 2015
NOVEL LABS INC	1.5MG	A202508	001	Feb 22, 2013
PERRIGO R AND D	1.5MG	A202334	001	Aug 20, 2014
XIROMED	1.5MG	A205329	001	Sep 18, 2018

OPCICON ONE-STEP

SUN PHARM	1.5MG	A202635	001	Sep 11, 2014
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PLAN B ONE-STEP

+! FDN CONSUMER	1.5MG	N021998	001	Jul 10, 2009
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LOPERAMIDE HYDROCHLORIDE

CAPSULE;ORAL

LOPERAMIDE HYDROCHLORIDE

+ BIONPHARMA INC	1MG	N021855	001	Aug 04, 2005
+!	2MG	N021855	002	Aug 04, 2005

SOLUTION;ORAL

IMODIUM A-D

+! J AND J CONSUMER INC	1MG/5ML	N019487	001	Mar 01, 1988
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LOPERAMIDE HYDROCHLORIDE

HI TECH PHARMA	1MG/5ML	A074352	001	Nov 17, 1995
PERRIGO	1MG/5ML	A073243	001	Jan 21, 1992
WOCKHARDT BIO AG	1MG/5ML	A074730	001	Aug 28, 1997

SUSPENSION;ORAL

IMODIUM A-D

+! J AND J CONSUMER INC	1MG/7.5ML	N019487	002	Jul 08, 2004
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LOPERAMIDE HYDROCHLORIDE

PERRIGO R AND D	1MG/7.5ML	A091292	001	May 20, 2011
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TABLET;ORAL

IMODIUM A-D

+! J AND J CONSUMER INC	2MG	N019860	001	Nov 22, 1989
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OTC DRUG PRODUCT LIST

LOPERAMIDE HYDROCHLORIDE

TABLET; ORAL

LOPERAMIDE HYDROCHLORIDE

AUROBINDO PHARMA LTD	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-SYMPTOM RELIEF

+! J AND J CONSUMER INC	2MG;125MG	N021140 001	Nov 30, 2000
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LOPERAMIDE HYDROCHLORIDE AND SIMETHICONE

PERRIGO R AND D	2MG;125MG	A209837 001	Sep 05, 2018
SUN PHARM INDS LTD	2MG;125MG	A077500 001	Sep 06, 2006

TABLET, CHEWABLE; ORAL

LOPERAMIDE HYDROCHLORIDE AND SIMETHICONE

! PERRIGO	2MG;125MG	A076029 001	Aug 30, 2002
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LORATADINE

CAPSULE; ORAL

CLARITIN

+! BAYER HEALTHCARE LLC	10MG	N021952 001	Jun 16, 2008
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LORATADINE

MARKSANS PHARMA	10MG	A206214 001	Sep 23, 2016
STRIDES PHARMA	10MG	A211926 001	Jan 15, 2020

SUSPENSION; ORAL

LORATADINE

+! TARO	1MG/ML	N021734 001	Oct 04, 2005
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SYRUP; ORAL

CLARITIN

+! BAYER HEALTHCARE LLC	1MG/ML	N020641 002	Nov 27, 2002
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LORATADINE

AUROBINDO PHARMA LTD	1MG/ML	A208931 001	Jun 29, 2018
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
TEVA	1MG/ML	A075505 001	Nov 07, 2003
WOCKHARDT BIO AG	1MG/ML	A075815 001	Aug 20, 2004

TABLET; ORAL

CLARITIN

+! BAYER HEALTHCARE LLC	10MG	N019658 002	Nov 27, 2002
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CLARITIN HIVES RELIEF

+! BAYER HEALTHCARE LLC	10MG	N019658 003	Nov 19, 2003
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LORATADINE

APOTEX INC	10MG	A076471 001	Feb 14, 2006
AUROBINDO PHARMA LTD	10MG	A208314 001	Apr 16, 2018
GRANULES INDIA LTD	10MG	A210722 001	Dec 23, 2019
GUARDIAN DRUG	10MG	A207569 001	Mar 12, 2019
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

TABLET, CHEWABLE; ORAL

CHILDREN'S CLARITIN

+! BAYER HEALTHCARE LLC	5MG	N021891 001	Aug 23, 2006
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CLARITIN

+ BAYER HEALTHCARE LLC	10MG	N021891 002	Nov 21, 2018
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LORATADINE

PERRIGO PHARMA INTL	5MG	A210033 001	Jun 12, 2019
SUN PHARM	5MG	A210088 001	Apr 16, 2018

OTC DRUG PRODUCT LISTLORATADINE

TABLET, ORALLY DISINTEGRATING;ORAL
ALAVERT

GLAXOSMITHKLINE	10MG	N021375	001	Dec 19, 2002
CLARITIN HIVES RELIEF REDITAB				
+! BAYER HEALTHCARE	10MG	N020704	003	Nov 19, 2003
LLC				
CLARITIN REDITABS				
+! BAYER HEALTHCARE	5MG	N021993	001	Dec 12, 2006
LLC				
+!	10MG	N020704	002	Nov 27, 2002
LORATADINE				
ACTAVIS LABS FL INC	10MG	A075990	001	Nov 03, 2003
AUROBINDO PHARMA	10MG	A208477	001	Apr 11, 2018
LTD				
GLAXOSMITHKLINE	10MG	A075822	001	Feb 10, 2003
PERRIGO PHARMA INTL	10MG	A076011	001	Sep 29, 2003
LORATADINE REDIDOSE				
SUN PHARM INDS LTD	10MG	A077153	001	Apr 11, 2007

LORATADINE; PSEUDOEPHEDRINE SULFATE

TABLET, EXTENDED RELEASE;ORAL

CLARITIN-D				
+! BAYER HEALTHCARE	5MG;120MG	N019670	002	Nov 27, 2002
LLC				
CLARITIN-D 24 HOUR				
+! BAYER HEALTHCARE	10MG;240MG	N020470	002	Nov 27, 2002
LLC				
LORATADINE AND PSEUDOEPHEDRINE SULFATE				
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
CREAM;VAGINAL				
MICONAZOLE 3				
TARO	4%	A076773	001	Mar 02, 2005
MICONAZOLE 7				
P AND L	2%	A074164	001	Mar 29, 1996
MICONAZOLE NITRATE				
ACP NIMBLE	2%	A074366	001	Feb 22, 1996
PERRIGO	2%	A074760	001	May 15, 1997
PERRIGO R AND D	4%	A091366	001	Jan 15, 2010
TARO PHARMS	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, SUPPOSITORY;TOPICAL, VAGINAL				
M-ZOLE 3 COMBINATION PACK				
P AND L	2%,200MG	A074926	001	Apr 16, 1999
MICONAZOLE NITRATE				
PERRIGO R AND D	2%,1.2GM	A079114	001	Jun 02, 2010
MICONAZOLE NITRATE COMBINATION PACK				
PERRIGO	2%,200MG	A075329	001	Apr 20, 1999
MONISTAT 1 COMBINATION PACK				
+! MEDTECH PRODUCTS	2%,1.2GM	N021308	001	Jun 29, 2001
MONISTAT 3 COMBINATION PACK				
+! MEDTECH PRODUCTS	2%,200MG	N020670	002	Apr 16, 1996

OTC DRUG PRODUCT LIST

4-15 (of 20)

MICONAZOLE NITRATE

CREAM, SUPPOSITORY;TOPICAL, VAGINAL

MONISTAT 7 COMBINATION PACK

+! MEDTECH PRODUCTS 2%,100MG

N020288 002 Apr 26, 1993

SUPPOSITORY;VAGINAL

MICONAZOLE NITRATE

ACP NIMBLE 100MG

A074414 001 Apr 30, 1997

P AND L 100MG

A073507 001 Nov 19, 1993

! PERRIGO 100MG

A074395 001 Mar 20, 1997

MONISTAT 7

+! MEDTECH PRODUCTS 100MG

N018520 002 Feb 15, 1991

MINOXIDIL

AEROSOL, FOAM;TOPICAL

MEN'S ROGAINE

+! JOHNSON AND JOHNSON 5%

N021812 001 Jan 20, 2006

MINOXIDIL

PERRIGO ISRAEL 5%

A091344 001 Apr 28, 2011

MINOXIDIL (FOR MEN)

P AND L 5%

A208092 001 Feb 17, 2017

TARO 5%

A209074 001 Dec 31, 2018

MINOXIDIL (FOR WOMEN)

P AND L 5%

A208092 002 Jul 27, 2017

TARO 5%

A209074 002 Apr 22, 2019

WOMEN'S ROGAINE

+! JOHNSON AND JOHNSON 5%

N021812 002 Feb 28, 2014

SOLUTION;TOPICAL

MINOXIDIL (FOR MEN)

HI TECH PHARMA 2%

A074731 001 Dec 24, 1996

L PERRIGO CO 2%

A075357 001 Jul 30, 1999

P AND L 2%

A074588 001 Apr 05, 1996

WOCKHARDT BIO AG 2%

A074767 001 Feb 28, 1997

MINOXIDIL (FOR WOMEN)

HI TECH PHARMA 2%

A074731 002 May 11, 2005

L PERRIGO CO 2%

A075357 002 Jul 30, 1999

MINOXIDIL EXTRA STRENGTH (FOR MEN)

AVACOR PRODS 5%

A075619 001 Nov 17, 2000

P AND L 5%

A075518 001 Nov 17, 2000

PERRIGO 5%

A075598 001 Jun 13, 2001

PERRIGO NEW YORK 5%

A075737 001 Mar 15, 2002

WOCKHARDT BIO AG 5%

A075438 001 Feb 27, 2003

ROGAINE (FOR MEN)

+! JOHNSON AND JOHNSON 2%

N019501 002 Feb 09, 1996

ROGAINE (FOR WOMEN)

+! JOHNSON AND JOHNSON 2%

N019501 003 Feb 09, 1996

ROGAINE EXTRA STRENGTH (FOR MEN)

+! JOHNSON AND JOHNSON 5%

N020834 001 Nov 14, 1997

THEROXIDIL

EI INC 2%

A078176 001 Nov 09, 2007

5%

A076239 001 Aug 24, 2004

NAPHAZOLINE HYDROCHLORIDE; PHENIRAMINE MALEATE

SOLUTION/DROPS;OPHTHALMIC

NAPHAZOLINE HYDROCHLORIDE AND PHENIRAMINE MALEATE

AKORN INC 0.025%;0.3%

A202795 001 Jan 24, 2013

ALTAIRE PHARMS INC 0.02675%;0.315%

A078208 001 Sep 27, 2010

NAPHCON-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

+! JOHNSON AND JOHNSON 0.025%;0.3%

N020485 001 Jan 31, 1996

NAPROXEN SODIUM

CAPSULE;ORAL

NAPROXEN SODIUM

+! BIONPHARMA INC EQ 200MG BASE

N021920 001 Feb 17, 2006

CATALENT EQ 200MG BASE

A202807 001 Jan 04, 2019

PURACAP PHARM LLC EQ 200MG BASE

A208363 001 Mar 15, 2018

TABLET;ORAL

ALEVE

+! BAYER 220MG

N020204 002 Jan 11, 1994

OTC DRUG PRODUCT LIST

4-16 (of 20)

NAPROXEN SODIUM

TABLET; ORAL

NAPROXEN SODIUM

AMNEAL PHARMS NY	220MG	A079096	001	Dec 16, 2008
AUROBINDO PHARMA LTD	220MG	A205497	001	Mar 18, 2016
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
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
SUN PHARM INDS LTD	220MG	A091183	001	May 20, 2011
YICHANG HUMANWELL	220MG	A212033	001	Aug 30, 2019

NAPROXEN SODIUM; PSEUDOEPHEDRINE HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL

ALEVE-D SINUS & COLD

+! BAYER	220MG;120MG	N021076	001	Nov 29, 1999
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NAPROXEN SODIUM AND PSEUDOEPHEDRINE HYDROCHLORIDE

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

+ SANOFI AVENTIS US	7MG/24HR	N020165	006	Aug 02, 1996
+	14MG/24HR	N020165	005	Aug 02, 1996
+!	21MG/24HR	N020165	004	Aug 02, 1996

NICOTINE

AVEVA	7MG/24HR	A074612	002	Jul 28, 2003
	14MG/24HR	A074612	003	Oct 20, 1997
	21MG/24HR	A074612	001	Oct 20, 1997

NICOTINE POLACRILEX

GUM, CHEWING; BUCCAL

NICORETTE

+ GLAXOSMITHKLINE	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)

+ GLAXOSMITHKLINE	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	A076776	001	Sep 16, 2004
	EQ 2MG BASE	A076777	001	Sep 16, 2004
	EQ 4MG BASE	A076778	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	A078967	001	Apr 23, 2008
	EQ 2MG BASE	A091349	001	Jul 20, 2011
	EQ 2MG BASE	A206394	001	Dec 15, 2016
	EQ 4MG BASE	A078326	001	Oct 30, 2006

OTC DRUG PRODUCT LIST

4-17 (of 20)

NICOTINE POLACRILEX

GUM, CHEWING;BUCCAL

NICOTINE POLACRILEX

EQ 4MG BASE	A078546	001	May 24, 2007
EQ 4MG BASE	A078968	001	Apr 23, 2008
EQ 4MG BASE	A091354	001	Jul 20, 2011
EQ 4MG BASE	A206393	001	Dec 15, 2016

TROCHE/LOZENGE;ORAL

NICORETTE

+ GLAXOSMITHKLINE CONS	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

DR REDDYS LABS SA

EQ 2MG BASE	A212796	001	Jan 08, 2020
EQ 2MG BASE	A212983	001	Feb 21, 2020
EQ 4MG BASE	A212796	002	Jan 08, 2020
EQ 4MG BASE	A212983	002	Feb 21, 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 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
EQ 2MG BASE	A077007	001	Jan 31, 2006
EQ 2MG BASE	A090711	001	Jul 10, 2009
EQ 2MG BASE	A090821	001	Jul 10, 2009
EQ 2MG BASE	A203690	001	Oct 09, 2012
EQ 4MG BASE	A077007	002	Jan 31, 2006
EQ 4MG BASE	A090711	002	Jul 10, 2009
EQ 4MG BASE	A090821	002	Jul 10, 2009
EQ 4MG BASE	A203690	002	Oct 09, 2012
EQ 2MG BASE	A207868	001	Feb 07, 2019
EQ 4MG BASE	A207868	002	Feb 07, 2019

PERRIGO R AND D

PLD ACQUISITIONS

NIZATIDINE

TABLET;ORAL

AXID AR

+	GLAXOSMITHKLINE	75MG	N020555	001	May 09, 1996
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NONOXYNOL-9

SPONGE;VAGINAL

TODAY

+	MAYER LABS INC	1GM	N018683	001	Apr 01, 1983
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OMEPRAZOLE

TABLET, DELAYED RELEASE;ORAL

OMEPRAZOLE

+	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 PHARMA	20MG	N209400	001	Jul 05, 2017
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OMEPRAZOLE MAGNESIUM

CAPSULE, DELAYED RELEASE;ORAL

OMEPRAZOLE MAGNESIUM

!	DR REDDYS LABS LTD	EQ 20MG BASE	A078878	001	Jun 05, 2009
	SPIL	EQ 20MG BASE	A210593	001	Jul 20, 2018

TABLET, DELAYED RELEASE;ORAL

OMEPRAZOLE MAGNESIUM

AUROBINDO PHARMA LTD	EQ 20MG BASE	A206877	001	Jun 06, 2018
PERRIGO R AND D	EQ 20MG BASE	A204152	001	Jul 30, 2015

OTC DRUG PRODUCT LIST

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OMEPRAZOLE MAGNESIUM

TABLET, DELAYED RELEASE;ORAL

PRILOSEC OTC

+! ASTRAZENECA PHARMS EQ 20MG BASE N021229 001 Jun 20, 2003

OMEPRAZOLE; SODIUM BICARBONATE

CAPSULE;ORAL

OMEPRAZOLE AND SODIUM BICARBONATE

ACTAVIS ELIZABETH 20MG;1.1GM A204137 001 Jul 15, 2016

AUROLIFE PHARMA LLC 20MG;1.1GM A204923 001 Nov 07, 2016

PAR PHARM 20MG;1.1GM A201946 001 Jul 15, 2016

PERRIGO R AND D 20MG;1.1GM A201361 001 Jul 15, 2016

ZYDUS PHARMS 20MG;1.1GM A203345 001 Mar 16, 2018

ZEGERID OTC

+! BAYER HEALTHCARE 20MG;1.1GM N022281 001 Dec 01, 2009
LLC

FOR SUSPENSION;ORAL

ZEGERID OTC

+! BAYER HEALTHCARE 20MG/PACKET;1.68GM/PACKET N022283 001 Jun 17, 2013
LLCORLISTAT

CAPSULE;ORAL

ALLI

+! GLAXOSMITHKLINE 60MG N021887 001 Feb 07, 2007
CONSOXYBUTYRIN

FILM, EXTENDED RELEASE;TRANSDERMAL

OXYTROL FOR WOMEN

+! ALLERGAN 3.9MG/24HR N202211 001 Jan 25, 2013

OXYMETAZOLINE HYDROCHLORIDE

SOLUTION/DROPS;OPHTHALMIC

VISINE L.R.

+! JOHNSON AND JOHNSON 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

GLYCOLAX

LANNETT CO INC 17GM/PACKET A090600 001 Oct 06, 2009

17GM/SCOOPFUL A090600 002 Oct 06, 2009

MIRALAX

+! BAYER HEALTHCARE 17GM/SCOOPFUL N022015 001 Oct 06, 2006
LLC

POLYETHYLENE GLYCOL 3350

ANI PHARMS INC 17GM/SCOOPFUL A202850 001 Dec 15, 2015

AUROBINDO PHARMA 17GM/SCOOPFUL A209017 001 Apr 09, 2018

LTD

MYLAN 17GM/PACKET A078915 001 Oct 06, 2009

17GM/SCOOPFUL A078915 002 Oct 06, 2009

NEXGEN PHARMA 17GM/SCOOPFUL A090812 001 Oct 07, 2009

NOVEL LABS INC 17GM/SCOOPFUL A091077 001 Oct 06, 2009

NOVELGENIX THERAPS 17GM/SCOOPFUL A202071 001 Dec 28, 2012

NUVO PHARMS INC 17GM/SCOOPFUL A206105 001 Oct 28, 2016

PAR PHARM 17GM/SCOOPFUL A079214 001 Jan 31, 2013

PERRIGO R AND D 17GM/PACKET A090685 001 Oct 06, 2009

17GM/SCOOPFUL A090685 002 Oct 06, 2009

STRIDES PHARMA 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

CO

THYROSHIELD

! ARCO PHARMS LLC 65MG/ML A077218 001 Jan 12, 2005

OTC DRUG PRODUCT LIST

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POTASSIUM IODIDE

TABLET; ORAL

IOSAT

+	ANBEX	65MG	N018664	002	May 12, 2011
+	!	130MG	N018664	001	Oct 14, 1982

THYROSAFE

!	RECIP	65MG	A076350	001	Sep 10, 2002
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POVIDONE-IODINE

SOLUTION; TOPICAL

POVIDONE IODINE

+	!	ALLEGIANCE HLTHCARE	1%	N019522	001	Mar 31, 1989
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SPONGE; TOPICAL

E-Z SCRUB 201

+	!	BECTON DICKINSON	20%	N019240	001	Nov 29, 1985
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E-Z SCRUB 241

+	!	BECTON DICKINSON	10%	N019476	001	Jan 07, 1987
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PSEUDOEPHEDRINE HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL

PSEUDOEPHEDRINE HYDROCHLORIDE

AUROBINDO PHARMA LTD	120MG	A209008	001	Jun 09, 2017
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LTD

L PERRIGO CO	120MG	A075153	001	Feb 26, 1999
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SUN PHARM INDS LTD

120MG	A077442	001	Sep 28, 2005
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SUDAFED 12 HOUR

!	MCNEIL CONS	120MG	A073585	001	Oct 31, 1991
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SUDAFED 24 HOUR

+	!	J AND J CONSUMER INC	240MG	N020021	002	Dec 15, 1992
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INC

PURIFIED WATER

SOLUTION; OPHTHALMIC

PUR-WASH

+	!	NIAGARA PHARMS	98.3%	N022305	001	Sep 01, 2011
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RANITIDINE HYDROCHLORIDE

TABLET; ORAL

RANITIDINE HYDROCHLORIDE

APOTEX INC	EQ 75MG BASE	A075167	001	May 04, 2000
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	EQ 150MG BASE	A200172	001	May 31, 2012
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AUROBINDO PHARMA LTD

EQ 75MG BASE	A207579	001	Nov 13, 2017
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LTD

EQ 150MG BASE

A207578	001	Nov 13, 2017
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DR REDDYS LABS LTD

EQ 75MG BASE

A075294	001	Mar 28, 2000
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EQ 150MG BASE

A078192	001	Aug 31, 2007
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GRANULES INDIA LTD

EQ 150MG BASE

A210243	001	Aug 20, 2018
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EQ 150MG BASE

A210243	002	Aug 20, 2018
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PERRIGO

EQ 75MG BASE

A076195	001	Aug 30, 2002
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PERRIGO R AND D

EQ 150MG BASE

A091429	001	May 11, 2011
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EQ 150MG BASE

A091429	002	May 11, 2011
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STRIDES PHARMA

EQ 75MG BASE

A201745	001	Feb 29, 2012
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EQ 75MG BASE

A209160	001	Mar 05, 2018
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EQ 150MG BASE

A200536	001	Jun 28, 2011
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EQ 150MG BASE

A209161	001	Feb 22, 2018
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UNIQUE PHARM LABS

EQ 75MG BASE

A210250	001	Aug 30, 2019
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EQ 150MG BASE

A210228	001	Aug 30, 2019
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EQ 75MG BASE

A076760	001	Feb 24, 2006
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ZANTAC 150

+	!	SANOFI US	EQ 150MG BASE	N021698	001	Aug 31, 2004
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+			EQ 150MG BASE	N021698	002	Mar 13, 2007
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ZANTAC 75

+		SANOFI US	EQ 75MG BASE	N020520	001	Dec 19, 1995
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SODIUM CHLORIDE

AEROSOL, METERED; INHALATION

BRONCHO SALINE

+	!	BLAIREX	0.9%	N019912	001	Sep 03, 1992
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TERBINAFINE

GEL; TOPICAL

LAMISIL AT

+	!	GLAXOSMITHKLINE CONS	1%	N021958	001	Jul 24, 2006
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OTC DRUG PRODUCT LISTTERBINAFINE HYDROCHLORIDE

CREAM; TOPICAL

LAMISIL

+! GLAXOSMITHKLINE 1%

N020980 001 Mar 09, 1999

TERBINAFINE HYDROCHLORIDE

TARO

1%

A077511 001 Jul 02, 2007

SOLUTION; TOPICAL

LAMISIL AT

+! GLAXOSMITHKLINE 1%

N021124 001 Mar 17, 2000

CONS

SPRAY; TOPICAL

LAMISIL AT

+! GLAXOSMITHKLINE 1%

N021124 002 Mar 17, 2000

CONS

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

+! SANOFI AVENTIS US 0.055MG/SPRAY

N020468 002 Oct 11, 2013

TRIAMCINOLONE ACETONIDE

PERRIGO ISRAEL 0.055MG/SPRAY

A078104 002 Nov 14, 2014

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ABACAIVIR SULFATE; LAMIVUDINE

TABLET; ORAL

ABACAIVIR SULFATE AND LAMIVUDINE

ZYDUS PHARMS EQ 600MG BASE;300MG A208990 001 Nov 15, 2018

ABARELIX

INJECTABLE; INTRAMUSCULAR

PLENAXIS

SPECIALITY EUROPEAN 100MG/VIAL N021320 001 Nov 25, 2003

ACAMPROSATE CALCIUM

TABLET, DELAYED RELEASE; ORAL

ACAMPROSATE CALCIUM

BARR LABS DIV TEVA 333MG A200143 001 Nov 18, 2013

CAMPRAL

+ FOREST LABS 333MG ** N021431 001 Jul 29, 2004

ACARBOSE

TABLET; ORAL

ACARBOSE

MYLAN 25MG A091053 001 Jan 06, 2011

50MG A091053 002 Jan 06, 2011

100MG A091053 003 Jan 06, 2011

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

ANI PHARMS INC EQ 200MG BASE A074007 001 Oct 18, 1995

EQ 400MG BASE A074007 002 Oct 18, 1995

SECTRAL

+ PROMIUS PHARMA EQ 200MG BASE N018917 001 Dec 28, 1984

+ EQ 400MG BASE N018917 003 Dec 28, 1984

ACETAMINOPHEN

INJECTABLE; INJECTION

INJECTAPAP

ORTHO MCNEIL PHARM 100MG/ML N017785 001 Mar 07, 1986

SUPPOSITORY; RECTAL

ACEPHEN

ACP NIMBLE 120MG A072218 001 Mar 27, 1992

325MG N018060 003 Dec 18, 1986

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

TYLENOL

J AND J CONSUMER INC 120MG N017756 002

650MG N017756 001

TABLET, EXTENDED RELEASE; ORAL

ACETAMINOPHEN

SUN PHARM INDS LTD 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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

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

TABLET; ORAL

BUTALBITAL AND ACETAMINOPHEN

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 PHARMS 500MG; 50MG; 40MG A040261 001 Oct 28, 1998

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

GILBERT LABS 325MG; 50MG; 40MG A087629 001 Nov 13, 1984

HIKMA PHARMS 325MG; 50MG; 40MG A089718 001 Jun 12, 1995

500MG; 50MG; 40MG A040336 001 Aug 18, 1999

MIKART 750MG; 50MG; 40MG A040496 001 Dec 23, 2003

MIRROR PHARMS LLC 500MG; 50MG; 40MG A040883 001 Dec 23, 2008

NOVAST LABS 325MG; 50MG; 40MG A040864 001 Dec 01, 2008

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

PHRENILIN WITH CAFFEINE AND CODEINE

VALEANT 325MG; 50MG; 40MG; 30MG A074911 001 Aug 22, 2001

DISCONTINUED DRUG PRODUCT LIST

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** 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

TABLET;ORAL

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
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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

ACI HEALTHCARE LTD	120MG/5ML;12MG/5ML	A086366	001	
ACTAVIS MID ATLANTIC	120MG/5ML;12MG/5ML	A085861	001	
DAVA PHARMS INC	120MG/5ML;12MG/5ML	A040098	001	Sep 20, 1996
LANNETT CO INC	120MG/5ML;12MG/5ML	A091238	001	Nov 10, 2011
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
DURAMED PHARMS BARR	300MG;15MG	A040223	001	Nov 18, 1997
	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	

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

ACETAMINOPHEN; CODEINE PHOSPHATE

TABLET; ORAL

ACETAMINOPHEN AND CODEINE PHOSPHATE

	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
PURACAP PHARM	300MG;30MG	A087762	001	Dec 10, 1982
PUREPAC PHARM	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	
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;15MG	A089997	001	Dec 28, 1994
	300MG;30MG	A087276	001	May 26, 1982
	300MG;30MG	A089998	001	Dec 28, 1994
	300MG;60MG	A087275	001	May 26, 1982
	300MG;60MG	A089999	001	Dec 28, 1994
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	
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	

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ACETAMINOPHEN; CODEINE PHOSPHATE

TABLET; ORAL

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	

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				
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
VINTAGE PHARMS	500MG/15ML;7.5MG/15ML	A040520	001	Oct 30, 2003
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				
ASCHER	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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

ACETAMINOPHEN; HYDROCODONE BITARTRATE

TABLET; ORAL

HYDROCODONE BITARTRATE AND ACETAMINOPHEN

	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 PINE BROOK	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 NY	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
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
	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	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
NOSTRUM LABS INC	325MG;2.5MG	A209924	001	Nov 16, 2018
	325MG;5MG	A209924	002	Nov 16, 2018

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

ACETAMINOPHEN; HYDROCODONE BITARTRATE

TABLET; ORAL

HYDROCODONE BITARTRATE AND ACETAMINOPHEN

	325MG; 7.5MG	A209924	003	Nov 16, 2018
	325MG; 10MG	A209924	004	Nov 16, 2018
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
SANDOZ	500MG; 5MG	A040149	001	Jan 27, 1997
	750MG; 7.5MG	A040149	002	Jan 27, 1997
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
	750MG; 7.5MG	A040281	002	Sep 30, 1998
WATSON LABS	325MG; 7.5MG	A040248	001	Apr 28, 2000
	325MG; 10MG	A040248	002	Apr 28, 2000
	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
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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

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
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TYLOX

JANSSEN PHARMS	500MG; 5MG	A088790	001	Dec 12, 1984
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TYLOX-325

ORTHO MCNEIL PHARM	325MG; 5MG	A088246	001	Nov 08, 1984
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SOLUTION; ORAL

OXYCODONE AND ACETAMINOPHEN

SPECGX LLC	325MG/5ML; 5MG/5ML	A040680	001	Sep 29, 2006
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OXYCODONE HYDROCHLORIDE AND ACETAMINOPHEN

VINTAGE PHARMS	325MG/5ML; 5MG/5ML	A203573	001	Dec 18, 2014
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ROXICET

HIKMA	325MG/5ML; 5MG/5ML	A089351	001	Dec 03, 1986
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TABLET; ORAL

OXYCODONE 2.5/APAP 500

BRISTOL MYERS SQUIBB	500MG; 2.5MG	A085910	001	
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OXYCODONE 5/APAP 500

BRISTOL MYERS SQUIBB	500MG; 5MG	A085911	001	
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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	
DURAMED PHARMS BARR	325MG; 5MG	A040272	001	Jun 30, 1998
MALLINCKRODT	500MG; 7.5MG	A040550	001	Jun 30, 2004
	650MG; 10MG	A040550	002	Jun 30, 2004
MAYNE PHARMA INC	500MG; 7.5MG	A090177	005	Oct 20, 2008
	650MG; 10MG	A090177	006	Oct 20, 2008
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
WATSON LABS	500MG; 7.5MG	A040371	001	Dec 29, 2000
	650MG; 10MG	A040371	002	Dec 29, 2000

PERCOCET

VINTAGE PHARMS LLC	325MG; 5MG	A085106	002	
	500MG; 7.5MG	A040341	001	Jul 26, 1999
	650MG; 10MG	A040341	002	Jul 26, 1999

ROXICET 5/500

ROXANE	500MG; 5MG	A089775	001	Jan 12, 1989
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TABLET, EXTENDED RELEASE; ORAL

XARTEMIS XR

+ MALLINCKRODT INC	325MG; 7.5MG	N204031	001	Mar 11, 2014
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ACETAMINOPHEN; OXYCODONE HYDROCHLORIDE; OXYCODONE TEREPHTHALATE

CAPSULE; ORAL

TYLOX

ORTHO MCNEIL PHARM	500MG; 4.5MG; 0.38MG	A085375	001	
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DISCONTINUED DRUG PRODUCT LIST

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** 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

APOTEX	325MG;37.5MG	A078778 001	Apr 07, 2014
CSPC OUYI	325MG;37.5MG	A076914 001	Jul 26, 2006
PAR PHARM	325MG;37.5MG	A076475 001	Apr 21, 2005

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

ACETAZOLAMIDE

CAPSULE, EXTENDED RELEASE;ORAL

ACETAZOLAMIDE

ACCORD HLTHCARE	500MG	A207659 001	Oct 18, 2018
MYLAN	500MG	A203917 001	Jun 18, 2019

DIAMOX

+ TEVA BRANDED PHARM	500MG	N012945 001	
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TABLET;ORAL

ACETAZOLAMIDE

ALRA	250MG	A083320 001	
ASCOT	250MG	A087686 001	Oct 20, 1982
HERITAGE PHARMA	250MG	A088882 001	Oct 22, 1985
SUN PHARM INDUSTRIES	125MG	A089753 002	Jun 22, 1988
	250MG	A089753 001	Jun 22, 1988
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

EMCURE PHARMS LTD	EQ 500MG BASE/VIAL	A202693 001	Dec 19, 2014
HOSPIRA	EQ 500MG BASE/VIAL	A040108 001	Oct 30, 1995
PAR STERILE PRODUCTS	EQ 500MG BASE/VIAL	A205358 001	Jun 20, 2017

DIAMOX

+ TEVA WOMENS	EQ 500MG BASE/VIAL **	N009388 001	
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ACETIC ACID, GLACIAL

SOLUTION/DROPS;OTIC

ACETASOL

ACTAVIS MID ATLANTIC	2%	A087146 001	
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ACETIC ACID

KV PHARM	2%	A085493 001	
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ORLEX

WARNER CHILCOTT	2%	A086845 001	
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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
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BOROFAIR

PHARMAFAIR	2%;0.79%	A088606 001	Aug 21, 1985
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DOMEBORO

BAYER PHARMS	2%;0.79%	A084476 001	
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ACETIC ACID, GLACIAL; DESONIDE

SOLUTION/DROPS;OTIC

TRIDESILON

BAYER PHARMS	2%;0.05%	N017914 001	
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ACETIC ACID, GLACIAL; HYDROCORTISONE

SOLUTION/DROPS;OTIC

ACETASOL HC

ACTAVIS MID ATLANTIC	2%;1%	A087143 001	Jan 13, 1982
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ACETIC ACID W/ HYDROCORTISONE

KV PHARM	2%;1%	A085492 001	
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HYDROCORTISONE AND ACETIC ACID

BAUSCH AND LOMB	2%;1%	A040097 001	Oct 31, 1994
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WOCKHARDT	2%;1%	A040168 001	Aug 30, 1996
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ORLEX HC

WARNER CHILCOTT	2%;1%	A086844 001	
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ACETIC ACID, GLACIAL; HYDROCORTISONE; NEOMYCIN SULFATE

SUSPENSION/DROPS;OTIC

NEO-CORT-DOME

BAYER PHARMS	2%;1%;EQ 0.35% BASE	N050238 001	
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DISCONTINUED DRUG PRODUCT LIST

6-11(of 430)

** See List Footnote

ACETOHEXAMIDE

TABLET; ORAL

ACETOHEXAMIDE

ANI PHARMS INC	250MG	A070869 001	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	
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ACETRIZOATE SODIUM

SOLUTION; INTRAUTERINE

SALPIX

ORTHO MCNEIL PHARM	53%	N009008 001	
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ACETYLCHOLINE CHLORIDE

FOR SOLUTION; OPHTHALMIC

MIOCHOL

+ NOVARTIS	20MG/VIAL **	N016211 001	
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ACETYLCYSTEINE

INJECTABLE; INTRAVENOUS

ACETYLCYSTEINE

MYLAN INSTITUTIONAL	6GM/30ML (200MG/ML)	A203624 001	Jun 19, 2015
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SOLUTION; INHALATION, ORAL

ACETYLCYSTEINE

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	
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+ MUCOSIL-10	20% **	N013601 001	
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DEY

DEY	10%	A070575 001	Oct 14, 1986
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MUCOSIL-20

DEY	20%	A070576 001	Oct 14, 1986
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TABLET, EFFERVESCENT; ORAL

CETYLEV

+ ARBOR PHARMS LLC	500MG	N207916 001	Jan 29, 2016
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+ 2.5GM		N207916 002	Jan 29, 2016
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ACETYLCYSTEINE; ISOPROTERENOL HYDROCHLORIDE

SOLUTION; INHALATION

MUCOMYST W/ ISOPROTERENOL

MEAD JOHNSON	10%; 0.05%	N017366 001	
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ACETYLDIGITOXIN

TABLET; ORAL

ACYLANID

NOVARTIS	0.1MG	N009436 001	
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ACITRETIN

CAPSULE; ORAL

ACITRETIN

MYLAN	17.5MG	A203707 001	Sep 10, 2015
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	22.5MG	A203707 002	Sep 10, 2015
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SORIATANE

+ STIEFEL LABS INC	17.5MG	N019821 003	Aug 06, 2009
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+ 22.5MG		N019821 004	Aug 06, 2009
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DISCONTINUED DRUG PRODUCT LIST

6-12(of 430)

** See List Footnote

ACRISORCIN

CREAM;TOPICAL

AKRINOL

SCHERING

2MG/GM

N012470 001

ACYCLOVIR

CAPSULE;ORAL

ACYCLOVIR

ACTAVIS ELIZABETH

200MG

A074906 001 Aug 26, 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

A074727 001 Apr 22, 1997

200MG

A074977 001 Apr 13, 1998

RANBAXY

200MG

A074975 001 Sep 30, 1998

ROXANE

200MG

A074570 002 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

+ MYLAN

200MG

N018828 001 Jan 25, 1985

OINTMENT;OPHTHALMIC

AVACLYR

+ FERA PHARMS LLC

3%

N202408 001 Mar 29, 2019

OINTMENT;TOPICAL

ACYCLOVIR

PERRIGO UK FINCO

5%

A205659 001 Feb 20, 2019

TABLET;ORAL

ACYCLOVIR

ACTAVIS ELIZABETH

400MG

A074870 001 Jun 05, 1997

800MG

A074870 002 Jun 05, 1997

CHARTWELL MOLECULES

400MG

A074834 001 Apr 24, 1997

800MG

A074834 002 Apr 24, 1997

IVAX SUB TEVA PHARMS

400MG

A074836 001 Apr 22, 1997

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

400MG

A075211 001 Sep 28, 1998

800MG

A074976 002 Apr 13, 1998

800MG

A075211 002 Sep 28, 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

+ MYLAN

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

ATHENEX INC

EQ 500MG BASE/VIAL

A074596 002 Apr 22, 1997

EQ 1GM BASE/VIAL

A074596 001 Apr 22, 1997

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 PHARMS

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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

6-13(of 430)

** See List Footnote

ACYCLOVIR SODIUMINJECTABLE; INJECTION
ACYCLOVIR SODIUM

	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
MYLAN LABS LTD	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
+		EQ 500MG BASE/VIAL **	N018603 001
+		EQ 1GM BASE/VIAL **	N018603 002

ADAPALENESOLUTION; TOPICAL
DIFFERIN

+	GALDERMA LABS LP	0.1% **	N020338 001	May 31, 1996
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ADAPALENE; BENZOYL PEROXIDE

GEL; TOPICAL

ADAPALENE AND BENZOYL PEROXIDE

	TARO	0.3%; 2.5%	A209148 001	Oct 17, 2018
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ADENOSINE

INJECTABLE; INJECTION

ADENOCARD

+	ASTELLAS	3MG/ML **	N019937 002	Oct 30, 1989
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ADENOSINE

	AM REGENT	3MG/ML	A090010 001	Apr 28, 2009
	TEVA PHARMS USA	3MG/ML	A076564 001	Jun 16, 2004
		3MG/ML	A078676 001	Jul 31, 2008
	WEST-WARD PHARMS INT	3MG/ML	A076501 001	Jun 16, 2004
	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

	EMCURE PHARMS LTD	60MG/20ML (3MG/ML)	A202313 001	Sep 15, 2014
		90MG/30ML (3MG/ML)	A202313 002	Sep 15, 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, CHEWABLE; ORAL

ALBENZA

	AMEDRA PHARMS LLC	200MG	N207844 001	Jun 11, 2015
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ALBUMIN CHROMATED CR-51 SERUM

INJECTABLE; INJECTION

CHROMALBIN

	ISO TEX	100uCi/VIAL	N017835 001
		250uCi/VIAL	N017835 002
		500uCi/VIAL	N017835 003

ALBUMIN IODINATED I-125 SERUM

INJECTABLE; INJECTION

RADIO-IODINATED (I 125) SERUM ALBUMIN (HUMAN)

	BAYER PHARMS	2.5uCi/AMP	N017846 001
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RADIOIODINATED SERUM ALBUMIN (HUMAN) IHSA I 125

	MALLINCKRODT	6.67uCi/ML	N017844 003
		10uCi/ML	N017844 001
		100uCi/ML	N017844 002

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

6-14(of 430)

** See List Footnote

ALBUMIN IODINATED I-131 SERUM

INJECTABLE; INJECTION

MEGATOPE

ISO TEX	2mCi/VIAL	N017837	003
	5uCi/AMP	N017837	004
	20uCi/AMP	N017837	005

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
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VENTOLIN

GLAXOSMITHKLINE	0.09MG/INH	N018473	001
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ALBUTEROL SULFATE

CAPSULE; INHALATION

VENTOLIN ROTACAPS

GLAXOSMITHKLINE	EQ 0.2MG BASE	N019489	001	May 04, 1988
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SOLUTION; INHALATION

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 AND LOMB	EQ 0.083% BASE	A075358	001	Mar 29, 2000
	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
HI TECH PHARMA	EQ 0.083% BASE	A075063	001	Feb 09, 1999
LANDELA PHARM	EQ 0.083% BASE	A077569	001	Apr 04, 2006
MYLAN SPECLT	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 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
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+	EQ 0.5% BASE **	N019243	001	Jan 14, 1987
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VENTOLIN

+ GLAXOSMITHKLINE	EQ 0.083% BASE **	N019773	001	Apr 23, 1992
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	EQ 0.5% BASE **	N019269	002	Jan 16, 1987
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SYRUP; ORAL

ALBUTEROL SULFATE

ACTAVIS MID ATLANTIC	EQ 2MG BASE/5ML	A075262	001	Mar 30, 1999
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
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VENTOLIN

GLAXOSMITHKLINE	EQ 2MG BASE/5ML **	N019621	001	Jun 10, 1987
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TABLET; ORAL

ALBUTEROL SULFATE

AM THERAP	EQ 2MG BASE	A072449	001	Dec 05, 1989
	EQ 4MG BASE	A072450	001	Dec 05, 1989
COPLEY PHARM	EQ 2MG BASE	A072966	001	Nov 22, 1991
	EQ 4MG BASE	A072967	001	Nov 22, 1991
DAVA PHARMS INC	EQ 2MG BASE	A072860	002	Dec 20, 1989
	EQ 4MG BASE	A072860	001	Dec 20, 1989
PLIVA	EQ 2MG BASE	A072316	001	Dec 05, 1989
	EQ 4MG BASE	A072317	001	Dec 05, 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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

ALBUTEROL SULFATE

TABLET; ORAL

ALBUTEROL SULFATE

	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
YAOPHARMA CO LTD	EQ 2MG BASE	A072151 001	Dec 05, 1989
	EQ 4MG BASE	A072152 001	Dec 05, 1989
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
TABLET, EXTENDED RELEASE; ORAL			
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			
DAVA PHARMS INC	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			
APOTEX INC	EQ 0.083% BASE; 0.017%	A077117 001	Dec 31, 2007
FOSUN PHARMA	EQ 0.083% BASE; 0.017%	A076867 001	Dec 21, 2006
TEVA PHARMS	EQ 0.083% BASE; 0.017%	A076724 001	Dec 31, 2007
DUONEB			
+ MYLAN SPECIALITY LP	EQ 0.083% BASE; 0.017% **	N020950 001	Mar 21, 2001

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

INJECTABLE; INJECTION

ALCOHOL 5% IN DEXTROSE 5%

MILES

5ML/100ML

A083483 001

ALCOHOL; DEXTROSE

INJECTABLE; INJECTION

ALCOHOL 10% AND DEXTROSE 5%

B BRAUN

10ML/100ML; 5GM/100ML

N004589 006

ALCOHOL 5% AND DEXTROSE 5%

B BRAUN

5ML/100ML; 5GM/100ML

N004589 004

ALCOHOL 5% IN D5-W

HOSPIRA

5ML/100ML; 5GM/100ML

A083263 001

ALCOHOL 5% IN DEXTROSE 5% IN WATER

BAXTER HLTHCARE

5ML/100ML; 5GM/100ML

A083256 001

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

ALENDRONATE SODIUM

SOLUTION; ORAL

FOSAMAX

+ MERCK

EQ 70MG BASE/75ML **

N021575 001 Sep 17, 2003

TABLET; ORAL

ALENDRONATE SODIUM

MYLAN

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 35MG BASE

A078638 001 Aug 04, 2008

EQ 70MG BASE

A076584 004 Aug 04, 2008

EQ 70MG BASE

A078638 002 Aug 04, 2008

TEVA PHARMS

EQ 35MG BASE

A076184 002 Aug 04, 2008

EQ 70MG BASE

A076184 001 Feb 06, 2008

UPSHER SMITH LABS

EQ 5MG BASE

A075871 001 Apr 22, 2009

EQ 10MG BASE

A075871 002 Apr 22, 2009

EQ 35MG BASE

A075871 004 Apr 22, 2009

EQ 40MG BASE

A075871 003 Apr 22, 2009

EQ 70MG BASE

A075871 005 Apr 22, 2009

FOSAMAX

+ MERCK AND CO INC

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

HERITAGE PHARMA

10MG

A079056 001 Jul 18, 2011

MYLAN

10MG

A079014 001 Jul 18, 2011

WOCKHARDT LTD

10MG

A090221 001 Aug 10, 2012

ALGLUCERASE

INJECTABLE; INJECTION

CEREDASE

GENZYME

10 UNITS/ML

N020057 004 May 08, 1992

80 UNITS/ML

N020057 003 Apr 05, 1991

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; 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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

6-17(of 430)

** See List Footnote

ALLOPURINOL

TABLET; ORAL

ALLOPURINOL

FOSUN PHARMA	100MG	A070268 001	Dec 31, 1985
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

ALPRAZOLAM

SOLUTION; ORAL

ALPRAZOLAM

ROXANE	0.5MG/5ML	A074314 001	Oct 31, 1993
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TABLET; ORAL

ALPRAZOLAM

ANI PHARMS INC	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
DAVA INTL INC	0.25MG	A074174 001	Oct 19, 1993
	0.5MG	A074174 002	Oct 19, 1993
	1MG	A074174 003	Oct 19, 1993
	2MG	A074174 004	Oct 19, 1993
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 PHARMS INC	0.25MG	A074046 001	Oct 19, 1993
	0.5MG	A074046 002	Oct 19, 1993
	1MG	A074046 003	Oct 19, 1993
	2MG	A074046 004	May 07, 1997
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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

ALPRAZOLAMTABLET, EXTENDED RELEASE;ORAL
ALPRAZOLAM

	3MG	A077198 004	May 13, 2010
ANI PHARMS INC	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
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
MYLAN	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
VINTAGE PHARMS	0.5MG	A078442 001	Oct 15, 2007
	1MG	A078442 002	Oct 15, 2007
	2MG	A078442 003	Oct 15, 2007
	3MG	A078442 004	Oct 15, 2007

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

ALPROSTADIL

INJECTABLE;INJECTION

CAVERJECT

PFIZER	0.005MG/ML	N020755 001	Oct 31, 1997	
	0.01MG/ML	N020755 002	Oct 01, 1997	
	0.02MG/ML	N020755 003	Oct 01, 1997	
+	PHARMACIA AND UPJOHN	0.005MG/VIAL	N020379 003	Jun 27, 1996
EDEX	AUXILIUM PHARMS LLC	0.005MG/VIAL	N020649 001	Jun 12, 1997

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
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DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

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				
+ SANOFI AVENTIS US	160MG;40MG	N018685	002	Dec 09, 1983

AMANTADINE HYDROCHLORIDE

CAPSULE; ORAL

AMANTADINE HYDROCHLORIDE

ACTAVIS ELIZABETH	100MG	A077659	001	Feb 23, 2006
LANNETT CO INC	100MG	A209221	001	Jun 15, 2017
WATSON LABS	100MG	A071382	001	Jan 21, 1987

SYMADINE

SOLVAY	100MG	A071000	001	Sep 04, 1986
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SYMMETREL

+ ENDO PHARMS	100MG **	N016020	001	
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SYRUP; ORAL

AMANTADINE HYDROCHLORIDE

G AND W LABS INC	50MG/5ML	A072655	001	Oct 30, 1990
LANNETT CO INC	50MG/5ML	A076352	001	Sep 10, 2004
TEVA PHARMS	50MG/5ML	A073115	001	Aug 23, 1991
VINTAGE	50MG/5ML	A077992	001	Dec 12, 2006

SYMMETREL

+ ENDO PHARMS	50MG/5ML **	N016023	002	
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TABLET; ORAL

SYMMETREL

+ ENDO PHARMS	100MG **	N018101	001	
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AMBENONIUM CHLORIDE

TABLET; ORAL

MYTELASE

SANOFI AVENTIS US	10MG	N010155	002	
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AMCINONIDE

CREAM; TOPICAL

CYCLOCORT

+ ASTELLAS	0.025%	N018116	001	
+	0.1%	N018116	002	

LOTION; TOPICAL

CYCLOCORT

+ ASTELLAS	0.1%	N019729	001	Jun 13, 1988
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OINTMENT; TOPICAL

CYCLOCORT

+ ASTELLAS	0.1%	N018498	001	
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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

ETHYOL

CLINIGEN	375MG/VIAL	N020221	002	Sep 10, 1999
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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
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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

AMIKACIN SULFATE

INJECTABLE; INJECTION

AMIKACIN SULFATE

	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
TEVA PHARMS USA	EQ 50MG BASE/ML	A064045 001	Sep 28, 1993
	EQ 250MG BASE/ML	A064045 002	Sep 28, 1993
WEST-WARD PHARMS INT	EQ 50MG BASE/ML	A063274 001	May 18, 1992
	EQ 250MG BASE/ML	A063275 001	May 18, 1992
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	
	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; HYDROCHLOROTHIAZIDE

TABLET; ORAL

AMILORIDE HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE

TEVA	EQ 5MG ANHYDROUS; 50MG	A070795 001	Apr 17, 1988
WATSON LABS	EQ 5MG ANHYDROUS; 50MG	A073334 001	Jul 19, 1991
YAOPHARMA CO LTD	EQ 5MG ANHYDROUS; 50MG	A073357 001	Nov 27, 1991
HYDRO-RIDE			
PAR PHARM	EQ 5MG ANHYDROUS; 50MG	A070347 001	Dec 25, 1990
MODURETIC 5-50			
+	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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

AMINO ACIDS

INJECTABLE; INJECTION

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
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 II 8.5%			
B BRAUN	8.5% (8.5GM/100ML)	N016822 002	
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
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 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 L; 22.4MG/100ML; 261MG/100ML; 205MG/100ML	N019714 002	Sep 12, 1988
HOSPIRA INC	4.25%; 36.8MG/100ML; 20GM/100ML; 51MG/100ML L; 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 L; 22.4MG/100ML; 261MG/100ML; 205MG/100ML	N019714 004	Sep 12, 1988
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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

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/100ML;22.4MG/100ML;104.5MG/100ML;205MG/100ML	N019712	002	Sep 08, 1988
HOSPIRA INC	4.25%;10GM/100ML;51MG/100ML;176.5MG/100ML;22.4MG/100ML;104.5MG/100ML;205MG/100ML	N019682	003	Nov 01, 1988

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;261MG/100ML;205MG/100ML	N019564	002	Dec 16, 1986
AMINOSYN II 4.25% W/ ELECTROLYTES IN DEXTROSE 25% IN PLASTIC CONTAINER				
ABBOTT	4.25%;25GM/100ML;51MG/100ML;22.4MG/100ML;261MG/100ML;205MG/100ML	N019564	004	Dec 16, 1986

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

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; 120MG/100ML; 49.3MG/100ML	N019564 001	Dec 16, 1986
	3.5%; 5GM/100ML; 30MG/100ML; 97MG/100ML; 120MG/100ML; 49.3MG/100ML	N019712 001	Sep 08, 1988
HOSPIRA INC	3.5%; 5GM/100ML; 30MG/100ML; 97MG/100ML; 120MG/100ML; 49.3MG/100ML	N019682 001	Nov 01, 1988

AMINOSYN II 4.25% M IN DEXTROSE 10% IN PLASTIC CONTAINER

ABBOTT	4.25%; 10GM/100ML; 30MG/100ML; 97MG/100ML; 120MG/100ML; 49.3MG/100ML	N019564 003	Dec 16, 1986
HOSPIRA INC	4.25%; 5GM/100ML; 30MG/100ML; 97MG/100ML; 120MG/100ML; 49.3MG/100ML	N019682 002	Nov 01, 1988

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; 216MG/100ML; 112MG/100ML	N020147 002	Oct 23, 1995
BAXTER HLTHCARE	2.75%; 15GM/100ML; 51MG/100ML; 261MG/100ML; 216MG/100ML; 112MG/100ML	N020147 003	Oct 23, 1995
BAXTER HLTHCARE	2.75%; 20GM/100ML; 51MG/100ML; 261MG/100ML; 216MG/100ML; 112MG/100ML	N020147 004	Oct 23, 1995
BAXTER HLTHCARE	2.75%; 25GM/100ML; 51MG/100ML; 261MG/100ML; 216MG/100ML; 112MG/100ML	N020147 005	Oct 23, 1995
BAXTER HLTHCARE	2.75%; 5GM/100ML; 51MG/100ML; 261MG/100ML; 216MG/100ML; 112MG/100ML	N020147 001	Oct 23, 1995
BAXTER HLTHCARE	4.25%; 10GM/100ML; 51MG/100ML; 261MG/100ML; 297MG/100ML; 77MG/100ML	N020147 007	Oct 23, 1995
BAXTER HLTHCARE	4.25%; 15GM/100ML; 51MG/100ML; 261MG/100ML; 297MG/100ML; 77MG/100ML	N020147 008	Oct 23, 1995
BAXTER HLTHCARE	4.25%; 20GM/100ML; 51MG/100ML; 261MG/100ML; 297MG/100ML; 77MG/100ML	N020147 009	Oct 23, 1995
BAXTER HLTHCARE	4.25%; 25GM/100ML; 51MG/100ML; 261MG/100ML; 297MG/100ML; 77MG/100ML	N020147 010	Oct 23, 1995
BAXTER HLTHCARE	4.25%; 5GM/100ML; 51MG/100ML; 261MG/100ML; 297MG/100ML; 77MG/100ML	N020147 006	Oct 23, 1995

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	
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; POTASSIUM ACETATE; SODIUM CHLORIDE; SODIUM PHOSPHATE, DIBASIC, HEPTAHYDRATE

INJECTABLE; INJECTION

AMINOSYN 3.5% M

ICU MEDICAL INC	3.5%; 21MG/100ML; 128MG/100ML; 234MG/100ML	N017789 005	
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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
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DISCONTINUED DRUG PRODUCT LIST

6-24(of 430)

** See List Footnote

<u>AMINO ACIDS; MAGNESIUM CHLORIDE; POTASSIUM ACETATE; POTASSIUM CHLORIDE; SODIUM ACETATE</u>					
INJECTABLE; INJECTION					
VEINAMINE 8%					
HOSPIRA INC	8%; 61MG/100ML; 211MG/100ML; 56MG/100ML; 38 8MG/100ML	N017957	001		
<u>AMINO ACIDS; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; POTASSIUM PHOSPHATE, DIBASIC; SODIUM CHLORIDE</u>					
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	
<u>AMINO ACIDS; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM CHLORIDE; SODIUM PHOSPHATE, DIBASIC</u>					
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	
<u>AMINO ACIDS; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM CHLORIDE; SODIUM PHOSPHATE, DIBASIC, HEPTAHYDRATE</u>					
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	
<u>AMINO ACIDS; MAGNESIUM CHLORIDE; POTASSIUM PHOSPHATE, DIBASIC; SODIUM ACETATE; SODIUM CHLORIDE</u>					
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	
TRAVASOL 5.5% W/ ELECTROLYTES					
BAXTER HLTHCARE	5.5%; 102MG/100ML; 522MG/100ML; 431MG/100M L; 224MG/100ML	N017493	001		
TRAVASOL 8.5% SULFITE FREE W/ ELECTROLYTES IN PLASTIC CONTAINER					
BAXTER HLTHCARE	8.5%; 102MG/100ML; 522MG/100ML; 594MG/100M L; 154MG/100ML	N020173	002	Oct 27, 1995	
TRAVASOL 8.5% W/ ELECTROLYTES					
BAXTER HLTHCARE	8.5%; 102MG/100ML; 522MG/100ML; 594MG/100M L; 154MG/100ML	N017493	002		
<u>AMINO ACIDS; MAGNESIUM CHLORIDE; POTASSIUM PHOSPHATE, DIBASIC; SODIUM CHLORIDE</u>					
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/100M L	N017673	005		
<u>AMINOCAPROIC ACID</u>					
INJECTABLE; INJECTION					
AMICAR					
+ CLOVER PHARMS	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	
SYRUP; ORAL					
AMINOCAPROIC ACID					
AKORN	1.25GM/5ML	A074759	001	Sep 02, 1998	

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

AMINOCAPROIC ACID

TABLET; ORAL

AMINOCAPROIC

AKORN

500MG

A075602 001 May 24, 2001

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

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

ACP NIMBLE

250MG

A085498 001 Mar 23, 1983

500MG

A085498 002 Jan 03, 1983

TABLET; ORAL

AMINOPHYLLIN

GD SEARLE LLC

100MG

N002386 002

200MG

N002386 003

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

AMINOPHYLLINE

TABLET; ORAL

AMINOPHYLLINE

ANI PHARMS INC	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
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	
LANNETT	100MG	A084588 001	
	200MG	A084588 002	
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	
	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	
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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	
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TEEBACIN

CONSOLIDATED MIDLAND	500MG	N007320 002	
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AMINOSALICYLATE SODIUM; AMINOSALICYLIC ACID

TABLET; ORAL

NEOPASALATE

MEDPOINTE PHARM HLC	846MG;112MG	A080059 002	
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DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

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
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AMIODARONE HYDROCHLORIDE

INJECTABLE; INJECTION

AMIODARONE HYDROCHLORIDE

AKORN	50MG/ML	A076232	001	Jul 05, 2006
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
HOSPIRA	50MG/ML	A075955	001	Oct 18, 2002
	50MG/ML	A076108	001	Oct 15, 2002
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
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NEXTERONE

+ BAXTER HLTHCARE	50MG/ML **	N022325	001	Dec 24, 2008
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TABLET; ORAL

AMIODARONE HYDROCHLORIDE

MYLAN	200MG	A075188	001	Feb 24, 1999
TEVA	200MG	A074895	001	Apr 16, 1999

CORDARONE

+ WYETH PHARMS	200MG **	N018972	001	Dec 24, 1985
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AMITRIPTYLINE HYDROCHLORIDE

CONCENTRATE; ORAL

ENDEP

ROCHE	40MG/ML	A085749	001
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INJECTABLE; INJECTION

AMITRIPTYLINE HYDROCHLORIDE

WATSON LABS	10MG/ML	A085594	001
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ELAVIL

ASTRAZENECA	10MG/ML	N012704	001
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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 INC	10MG	A085031	002	
	25MG	A085031	001	
	50MG	A085031	003	
	75MG	A085031	004	
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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

AMITRIPTYLINE HYDROCHLORIDE

TABLET;ORAL

AMITRIPTYLINE HYDROCHLORIDE

	150MG	A088426 001	Apr 30, 1984
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	
	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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

AMITRIPTYLINE HYDROCHLORIDE

TABLET;ORAL

AMITRIPTYLINE HYDROCHLORIDE

	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
	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

FRONTIDA BIOPHARM	EQ 12.5MG BASE;5MG	A070765	001	Dec 10, 1986
	EQ 25MG BASE;10MG	A070766	001	Dec 10, 1986
HERITAGE PHARMA	EQ 12.5MG BASE;5MG	A072052	001	Dec 16, 1988
	EQ 25MG BASE;10MG	A072053	001	Dec 16, 1988
PAR PHARM	EQ 12.5MG BASE;5MG	A072277	001	May 09, 1988
	EQ 25MG BASE;10MG	A072278	001	May 09, 1988
USL PHARMA	EQ 12.5MG BASE;5MG	A070477	001	Jan 12, 1988
	EQ 25MG BASE;10MG	A070478	001	Jan 12, 1988
LIMBITROL				
+	HERITAGE PHARMS INC	EQ 12.5MG BASE;5MG	**	N016949 001
LIMBITROL DS				
+	HERITAGE PHARMS INC	EQ 25MG BASE;10MG	**	N016949 002

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

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

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

HERITAGE PHARMA 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

IVAX SUB TEVA PHARMS 10MG; 2MG A070935 001 Sep 11, 1986

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

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

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
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
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
MYLAN	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
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 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 LAKE	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
TEVA	EQ 2.5MG BASE	A076846 001	Jun 28, 2007
	EQ 5MG BASE	A076846 002	Jun 28, 2007
	EQ 10MG BASE	A076846 003	Jun 28, 2007
YAOPHARMA CO LTD	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

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; BENAZEPRIL HYDROCHLORIDE

CAPSULE;ORAL

AMLODIPINE BESYLATE AND BENAZEPRIL HYDROCHLORIDE

CIPLA	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
MYLAN	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
	EQ 10MG BASE;40MG	A079047 002	Jul 05, 2011
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

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

AMLODIPINE BESYLATE; HYDROCHLOROTHIAZIDE; OLMESARTAN MEDOXOMIL

TABLET; ORAL

OLMESARTAN MEDOXOMIL, AMLODIPINE AND HYDROCHLOROTHIAZIDE

TEVA PHARMS USA	EQ 5MG BASE;12.5MG;20MG	A202491 001	Nov 03, 2016
	EQ 5MG BASE;12.5MG;40MG	A202491 002	Nov 03, 2016
	EQ 5MG BASE;25MG;40MG	A202491 003	Nov 03, 2016
	EQ 10MG BASE;12.5MG;40MG	A202491 004	Nov 03, 2016
	EQ 10MG BASE;25MG;40MG	A202491 005	Nov 03, 2016

AMLODIPINE BESYLATE; HYDROCHLOROTHIAZIDE; VALSARTAN

TABLET; ORAL

AMLODIPINE BESYLATE, VALSARTAN AND HYDROCHLOROTHIAZIDE

TEVA PHARMS	5MG;12.5MG;160MG	A200435 001	Sep 25, 2012
	5MG;25MG;160MG	A200435 002	Sep 25, 2012
	10MG;12.5MG;160MG	A200435 005	Sep 25, 2012
	10MG;25MG;160MG	A200435 003	Sep 25, 2012
	10MG;25MG;320MG	A200435 004	Sep 25, 2012
TORRENT	5MG;12.5MG;160MG	A201593 001	Jun 03, 2015
	5MG;25MG;160MG	A201593 002	Jun 03, 2015
	10MG;12.5MG;160MG	A201593 003	Jun 03, 2015
	10MG;25MG;160MG	A201593 004	Jun 03, 2015
	10MG;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
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

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
UNIV TX MD ANDERSON	30mCi-300mCi/8ML (3.75-37.5mCi/ML)	A203933 001	Jun 27, 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

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

AMMONIUM LACTATE

CREAM;TOPICAL

LAC-HYDRIN

SUN PHARM INDS INC EQ 12% BASE **

N020508 001 Aug 29, 1996

LOTION;TOPICAL

LAC-HYDRIN

+ SUN PHARM INDS INC EQ 12% BASE **

N019155 001 Apr 24, 1985

AMODIAQUINE HYDROCHLORIDE

TABLET;ORAL

CAMOQUIN HYDROCHLORIDE

PARKE DAVIS EQ 200MG BASE

N006441 001

AMOXAPINE

TABLET;ORAL

AMOXAPINE

UPSHER SMITH LABS

25MG

A072943 001 Jun 28, 1991

50MG

A072944 001 Jun 28, 1991

100MG

A072878 001 Jun 28, 1991

150MG

A072879 001 Jun 28, 1991

WATSON PHARMS TEVA

25MG

A072418 001 Aug 01, 1989

50MG

A072419 001 Aug 01, 1989

100MG

A072420 001 May 11, 1989

150MG

A072421 001 May 11, 1989

ASENDIN

LEDERLE

25MG

N018021 001

50MG

N018021 002

100MG

N018021 003

150MG

N018021 004

AMOXICILLIN

CAPSULE;ORAL

AMOXICILLIN

LABS ATRAL

250MG

A062528 001 Aug 07, 1985

500MG

A062528 002 Aug 07, 1985

MYLAN

250MG

A062067 001

500MG

A062067 002

SUN PHARM INDS LTD

250MG

A065016 001 Apr 08, 1999

500MG

A065016 002 Apr 08, 1999

TEVA

250MG

A062853 001 Dec 22, 1987

250MG

A063030 001 Feb 28, 1989

500MG

A062854 001 Dec 22, 1987

500MG

A063031 001 Feb 28, 1989

AMOXIL

+ GLAXOSMITHKLINE

250MG **

N050459 001

+

500MG **

N050459 002

TRIMOX

APOTHECON

250MG

A061885 001

250MG

A062098 001

250MG

A062152 001

250MG

A063099 001 Mar 20, 1992

500MG

A061885 002

500MG

A062098 002

500MG

A062152 002

500MG

A063099 002 Mar 20, 1992

UTIMOX

PARKE DAVIS

250MG

A062107 001

500MG

A062107 002

WYMOX

WYETH AYERST

250MG

A062120 001

500MG

A062120 002

FOR SUSPENSION;ORAL

AMOXICILLIN

AM ANTIBIOTICS

125MG/5ML

A062059 001

250MG/5ML

A062059 002

MYLAN

125MG/5ML

A062090 001

250MG/5ML

A062090 002

SUN PHARM INDS LTD

200MG/5ML

A065113 001 Nov 29, 2002

400MG/5ML

A065113 002 Nov 29, 2002

TEVA

125MG/5ML

A061931 001

125MG/5ML

A062946 001 Nov 01, 1988

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

AMOXICILLIN

FOR SUSPENSION;ORAL

AMOXICILLIN	250MG/5ML	A063001 001	Jan 06, 1989
AMOXIL			
+ GLAXOSMITHKLINE	50MG/ML **	N050460 005	
+	125MG/5ML **	N050460 001	
+	250MG/5ML **	N050460 002	
+ NEOPHARMA	200MG/5ML **	N050760 001	Apr 15, 1999
+	400MG/5ML **	N050760 002	Apr 15, 1999
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			
DAVA PHARMS INC	875MG	A065344 001	Jan 15, 2009
SUN PHARM INDS LTD	500MG	A065059 001	Nov 24, 2000
	875MG	A065059 002	Nov 24, 2000
AMOXIL			
+ NEOPHARMA	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
DAVA PHARMS INC	125MG	A064139 001	Jan 29, 1996
	250MG	A064139 002	Jan 29, 1996
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			
+ NEOPHARMA	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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

AMOXICILLIN; CLARITHROMYCIN; LANSOPRAZOLE

CAPSULE, CAPSULE, DELAYED REL PELLETS, TABLET;ORAL

LANSOPRAZOLE, AMOXICILLIN AND CLARITHROMYCIN

ANI PHARMS INC	500MG,N/A,N/A;N/A,500MG,N/A;N/A,N/A,30MG	A200218	001	Aug 30, 2013
PREVPAC				
+ TAKEDA PHARMS USA	500MG,N/A,N/A;N/A,500MG,N/A;N/A,N/A,30MG	N050757	001	Dec 02, 1997

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'				
+ NEOPHARMA	200MG/5ML;EQ 28.5MG BASE/5ML	N050725	001	May 31, 1996
AUGMENTIN '400'				
+ NEOPHARMA	400MG/5ML;EQ 57MG BASE/5ML	N050725	002	May 31, 1996
AUGMENTIN ES-600				
+ NEOPHARMA	600MG/5ML;EQ 42.9MG BASE/5ML	N050755	001	Jun 22, 2001

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
SUN PHARM INDS LTD	500MG;EQ 125MG BASE	A065109	001	Nov 04, 2002
	875MG;EQ 125MG BASE	A065102	001	Sep 17, 2002
AUGMENTIN '250'				
+ NEOPHARMA	250MG;EQ 125MG BASE **	N050564	001	Aug 06, 1984
AUGMENTIN '500'				
+ NEOPHARMA	500MG;EQ 125MG BASE **	N050564	002	Aug 06, 1984

TABLET, CHEWABLE;ORAL

AMOXICILLIN AND CLAVULANATE POTASSIUM

SANDOZ	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'				
+ NEOPHARMA	125MG;EQ 31.25MG BASE **	N050597	001	Jul 22, 1985
AUGMENTIN '200'				
+ NEOPHARMA	200MG;EQ 28.5MG BASE	N050726	001	May 31, 1996
AUGMENTIN '250'				
+ NEOPHARMA	250MG;EQ 62.5MG BASE **	N050597	002	Jul 22, 1985
AUGMENTIN '400'				
+ NEOPHARMA	400MG;EQ 57MG BASE	N050726	002	May 31, 1996

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

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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATE

CAPSULE, EXTENDED RELEASE; ORAL

DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE

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

TABLET; ORAL

ADDERALL 10

+ TEVA WOMENS	2.5MG;2.5MG;2.5MG;2.5MG **	N011522 007	Feb 13, 1996
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ADDERALL 12.5

+ TEVA WOMENS	3.125MG;3.125MG;3.125MG;3.125MG **	N011522 012	Aug 31, 2000
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ADDERALL 15

+ TEVA WOMENS	3.75MG;3.75MG;3.75MG;3.75MG **	N011522 013	Aug 31, 2000
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ADDERALL 20

+ TEVA WOMENS	5MG;5MG;5MG;5MG **	N011522 008	Feb 13, 1996
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ADDERALL 30

+ TEVA WOMENS	7.5MG;7.5MG;7.5MG;7.5MG **	N011522 010	May 12, 1997
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ADDERALL 5

+ TEVA WOMENS	1.25MG;1.25MG;1.25MG;1.25MG **	N011522 009	May 12, 1997
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ADDERALL 7.5

+ TEVA WOMENS	1.875MG;1.875MG;1.875MG;1.875MG **	N011522 011	Aug 31, 2000
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DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE

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
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	
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BIPHETAMINE 20

UCB INC	EQ 10MG BASE;EQ 10MG BASE	N010093 003	
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BIPHETAMINE 7.5

UCB INC	EQ 3.75MG BASE;EQ 3.75MG BASE	N010093 009	
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AMPHETAMINE SULFATE

TABLET; ORAL

AMPHETAMINE SULFATE

LANNETT	5MG	A083901 001	Aug 31, 1984
	10MG	A083901 002	Aug 31, 1984

AMPHOTERICIN B

CREAM; TOPICAL

FUNGIZONE

APOTHECON	3%	N050314 001	
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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	
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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	
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Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

AMPHOTERICIN B

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
AUROBINDO PHARMA	EQ 125MG BASE/VIAL	A065499 001	Aug 17, 2010
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	
HANFORD GC	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
HOSPIRA INC	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
	EQ 10GM BASE/VIAL	A202865 001	Sep 04, 2015
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
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
WEST-WARD PHARMS INT	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
	EQ 10GM BASE/VIAL	A062692 006	Jun 24, 1986
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	

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

AMPICILLIN SODIUM

INJECTABLE; INJECTION

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

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

PHARM ASSOC

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 500MG BASE/VIAL;EQ 250MG BASE/VIAL	N050608 003	Dec 31, 1986
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AMPICILLIN/AMPICILLIN TRIHYDRATE

CAPSULE; ORAL

AMCILL

PARKE DAVIS

EQ 250MG BASE	A062041 001
EQ 500MG BASE	A062041 002

AMPICILLIN TRIHYDRATE

AM ANTIBIOTICS

EQ 250MG BASE	A061602 001
EQ 500MG BASE	A061602 002

HERITAGE PHARMA

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

MYLAN

EQ 250MG BASE	A061755 001
EQ 500MG BASE	A061755 002

PUREPAC PHARM

EQ 250MG BASE	A061853 001
EQ 500MG BASE	A061853 002

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
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Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

AMPICILLIN/AMPICILLIN TRIHYDRATE

CAPSULE; ORAL

PRINCIPEN '250'	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		
AM ANTIBIOTICS	EQ 125MG BASE/5ML	A061601 001
	EQ 250MG BASE/5ML	A061601 002
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
	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

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

AMPICILLIN/AMPICILLIN TRIHYDRATE; PROBENECID

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

ACP NIMBLE

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

+ SHIRE LLC

EQ 1MG BASE **

N020333 002 Mar 14, 1997

ANAGRELIDE HYDROCHLORIDE

MYLAN

EQ 0.5MG BASE

A076811 001 Apr 18, 2005

EQ 1MG BASE

A076811 002 Apr 18, 2005

MYLAN PHARMS INC

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

UPSHER SMITH LABS

EQ 0.5MG BASE

A076683 001 Apr 18, 2005

EQ 1MG BASE

A076683 002 Apr 18, 2005

WATSON LABS

EQ 0.5MG BASE

A076417 001 Apr 18, 2005

EQ 1MG BASE

A076417 002 Apr 18, 2005

ANASTROZOLE

TABLET; ORAL

ANASTROZOLE

APOTEX INC

1MG

A200654 001 May 11, 2012

IMPAX LABS INC

1MG

A091242 001 May 31, 2012

LANNETT CO INC

1MG

A091331 001 Jan 05, 2011

MYLAN

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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

ANTAZOLINE PHOSPHATE; NAPHAZOLINE HYDROCHLORIDE

SOLUTION/DROPS;OPHTHALMIC

VASOCON-A

NOVARTIS

0.5%;0.05%

N018746 002 Jul 11, 1994

APIXABAN

TABLET;ORAL

APIXABAN

MYLAN

2.5MG

A210128 001 Dec 23, 2019

5MG

A210128 002 Dec 23, 2019

APOMORPHINE HYDROCHLORIDE

INJECTABLE;SUBCUTANEOUS

APOKYN

US WORLDMEDS

20MG/2ML (10MG/ML)

N021264 001 Apr 20, 2004

APREPITANT

CAPSULE;ORAL

EMEND

+ MERCK

40MG

N021549 003 Jun 30, 2006

APROTININ

INJECTABLE;INJECTION

TRASYLOL

BAYER HLTHCARE

10,000KIU/ML

N020304 001 Dec 29, 1993

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

ARGATROBAN

SOLUTION;INTRAVENOUS

ARGATROBAN IN DEXTROSE

SANDOZ

125MG/125ML (1MG/ML)

N201743 001 May 09, 2011

ARIPIRAZOLE

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

ARIPIRAZOLE

MYLAN

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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

ARIPIPIRAZOLE

TABLET, ORALLY DISINTEGRATING;ORAL

ABILIFY

+	20MG **	N021729 004	Jun 07, 2006
+	30MG **	N021729 005	Jun 07, 2006

ARMODAFINIL

TABLET;ORAL

ARMODAFINIL

APOTEX INC	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
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ARSENIC TRIOXIDE

INJECTABLE;INJECTION

TRISENOX

+	CEPHALON	1MG/ML **	N021248 001	Sep 25, 2000
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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
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ULTACAN FORTE

HANSAMED INC	4%;EQ 0.017MG BASE/1.7ML (4%; EQ 0.01MG BASE/ML)	A201750 001	Jul 11, 2017
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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;100IU/ML;0.2MG/ML;20MG/ML;2MG/ML;1.8MG/ML;	N006071 003	Oct 10, 1985
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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;20IU/ML;0.04MG/ML;4MG/ML;0.4MG/ML;0.36MG/	N018440 002	Aug 08, 1985
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M.V.I.-12 ADULT

HOSPIRA	10MG/ML;0.006MG/ML;0.5MCG/ML;1.5MG/ML;20IU/ML;0.04MG/ML;4MG/ML;0.4MG/ML;0.36MG/	N008809 004	Aug 08, 1985
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+

	20MG/ML;0.006MG/ML;0.05MCG/ML;1.5MG/ML;0.0005MG/ML;0.06MG/ML;4MG/ML;0.6MG/ML;0.36MG/ML;0.6MG/ML;0.1MG/ML;1MG/ML	N008809 006	Sep 09, 2004
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MVC PLUS

WATSON LABS	10MG/ML;0.006MG/ML;0.5MCG/ML;1.5MG/ML;20IU/ML;0.04MG/ML;4MG/ML;0.4MG/ML;0.36MG/	N018439 002	Aug 08, 1985
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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;20IU/ML;0.6MG/ML;4MG/ML;0.4MG/ML;0.36MG/M	N008809 005	Apr 22, 2004
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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 PHARMA INC	100MG/VIAL;0.06MG/VIAL;0.005MG/VIAL;15MG/VIAL;5MCG/VIAL;0.4MG/VIAL;40MG/VIAL;4MG/VIAL;3.6MG/VIAL;3MG/VIAL;1MG/VIAL;10MG/VIAL	N018933 002	Aug 08, 1985
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Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** 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

HOSPIRA	N/A, 80MG/VIAL; N/A, 0.02MG/VIAL; N/A, 0.001 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/VIAL; N/A, 1.4MG/VIAL; N/A, 1.2MG/VIAL; EQ 2,300 UNITS BASE/10ML, N/A; 7 IU/10ML, N/A	N020176 001	Dec 29, 1993
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ASPIRIN

TABLET; ORAL

BAYER EXTRA STRENGTH ASPIRIN FOR MIGRAINE PAIN

BAYER	500MG	N021317 001	Oct 18, 2001
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TABLET, EXTENDED RELEASE; ORAL

8-HOUR BAYER

BAYER	650MG	N016030 001	
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MEASURIN

BAYER	650MG	N016030 002	
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ASPIRIN; BUTALBITAL

TABLET; ORAL

AXOTAL

SAVAGE LABS	650MG; 50MG	A088305 001	Oct 13, 1983
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ASPIRIN; BUTALBITAL; CAFFEINE

CAPSULE; ORAL

BUTALBITAL, ASPIRIN AND CAFFEINE

NOSTRUM LABS INC	325MG; 50MG; 40MG	A078149 001	Jun 13, 2007
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WATSON LABS	325MG; 50MG; 40MG	A086231 002	Feb 12, 1985
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TABLET; ORAL

BUTALBITAL, ASPIRIN AND CAFFEINE

ACTAVIS ELIZABETH	325MG; 50MG; 40MG	A086710 002	Aug 23, 1983
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FOSUN PHARMA	325MG; 50MG; 40MG	A086398 002	Apr 06, 1984
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HALSEY	325MG; 50MG; 40MG	A089448 001	Dec 01, 1986
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IVAX PHARMS	325MG; 50MG; 40MG	A085441 002	Oct 31, 1984
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PURACAP PHARM	325MG; 50MG; 40MG	A087048 002	Dec 09, 1983
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QUANTUM PHARMICS	325MG; 50MG; 40MG	A088972 001	Jun 18, 1985
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WATSON LABS	325MG; 50MG; 40MG	A086237 002	Mar 23, 1984
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FIORINAL

+ ALLERGAN	325MG; 50MG; 40MG **	N017534 003	Apr 16, 1986
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LANORINAL

LANNETT	325MG; 50MG; 40MG	A086986 002	Oct 18, 1985
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ASPIRIN; BUTALBITAL; CAFFEINE; CODEINE PHOSPHATE

CAPSULE; ORAL

BUTALBITAL, ASPIRIN, CAFFEINE, AND CODEINE PHOSPHATE

VINTAGE PHARMS LLC	325MG; 50MG; 40MG; 30MG	A075351 001	Mar 05, 1999
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WATSON LABS	325MG; 50MG; 40MG; 30MG	A074359 001	Aug 31, 1995
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ASPIRIN; CAFFEINE; DIHYDROCODEINE BITARTRATE

CAPSULE; ORAL

SYNALGOS-DC

+ SUN PHARM INDUSTRIES	356.4MG; 30MG; 16MG	N011483 004	Sep 06, 1983
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ASPIRIN; CAFFEINE; ORPHENADRINE CITRATE

TABLET; ORAL

INVAGESIC

SANDOZ	385MG; 30MG; 25MG	A074817 001	Nov 27, 1996
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INVAGESIC FORTE

SANDOZ	770MG; 60MG; 50MG	A074817 002	Nov 27, 1996
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NORGESIC

+ MEDICIS	385MG; 30MG; 25MG **	N013416 003	Oct 27, 1982
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NORGESIC FORTE

+ MEDICIS	770MG; 60MG; 50MG **	N013416 004	Oct 27, 1982
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ORPHENADRINE CITRATE, ASPIRIN, AND CAFFEINE

SANDOZ	385MG; 30MG; 25MG	A074654 001	Dec 31, 1996
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	770MG; 60MG; 50MG	A074654 002	Dec 31, 1996
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STEVENS J	385MG; 30MG; 25MG	A074988 001	Apr 30, 1999
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	770MG; 60MG; 50MG	A074988 002	Apr 30, 1999
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Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

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; 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

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

OXFORD PHARMS 325MG; 200MG; 16MG

A040283 001 Dec 29, 1998

SOMA COMPOUND W/ CODEINE

MEDA PHARMS 325MG; 200MG; 16MG **

N012366 002 Jul 11, 1983

ASPIRIN; DIPYRIDAMOLE

CAPSULE, EXTENDED RELEASE; ORAL

ASPIRIN AND DIPYRIDAMOLE

ANI PHARMS INC 25MG; 200MG

A206964 001 Jan 18, 2017

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

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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

ASPIRIN; METHOCARBAMOL

TABLET; ORAL

ROBAXISAL

ROBINS AH

325MG; 400MG

N012281 001

ASPIRIN; OXYCODONE HYDROCHLORIDE; OXYCODONE TEREPHTHALATE

TABLET; ORAL

CODOXY

HALSEY

325MG; 4.5MG; 0.38MG

A087464 001 Jul 01, 1982

OXYCODONE AND ASPIRIN

ANI PHARMS INC

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 PHARMS

325MG; 4.5MG; 0.38MG **

N007337 006

PERCODAN-DEMI

ENDO PHARMS

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, TABLET; ORAL

PRAVIGARD PAC (COPACKAGED)

BRISTOL MYERS SQUIBB

325MG, N/A; N/A, 80MG

N021387 006 Jun 24, 2003

TABLET, TABLET, TABLET; ORAL

PRAVIGARD PAC (COPACKAGED)

BRISTOL MYERS SQUIBB

81MG, N/A; N/A, 20MG

N021387 001 Jun 24, 2003

81MG, N/A; N/A, 40MG

N021387 002 Jun 24, 2003

81MG, N/A; N/A, 80MG

N021387 003 Jun 24, 2003

325MG, N/A; N/A, 20MG

N021387 004 Jun 24, 2003

325MG, N/A; N/A, 40MG

N021387 005 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

ATAZANAVIR SULFATE

CAPSULE; ORAL

ATAZANAVIR SULFATE

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

REYATAZ

+ BRISTOL MYERS SQUIBB

EQ 100MG BASE **

N021567 001 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

DAVA PHARMS INC

25MG

A074099 001 Apr 28, 1992

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

ATENOLOL

TABLET; ORAL

ATENOLOL

INVATECH	25MG	A074265 001	Feb 28, 1994
	50MG	A074265 002	Feb 28, 1994
	100MG	A074265 003	Feb 28, 1994
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

ATENOLOL; CHLORTHALIDONE

TABLET; ORAL

ATENOLOL AND CHLORTHALIDONE

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	5MG	N021411 001	Nov 26, 2002
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ATORVASTATIN CALCIUM

TABLET; ORAL

ATORVASTATIN CALCIUM

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

+	MERCK SHARP DOHME	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
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DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

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
MYLAN LABS LTD	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

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
MYLAN LABS LTD	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

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
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SOLUTION; INTRAVENOUS, INTRAMUSCULAR, SUBCUTANEOUS, ENDOTRACHEAL

ATROPINE SULFATE ANSYR PLASTIC SYRINGE

+ HOSPIRA	0.5MG/5ML (0.1MG/ML) **	N021146	001	Jul 09, 2001
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ATROPINE SULFATE; DIFENOXIN HYDROCHLORIDE

TABLET; ORAL

MOTOFEN HALF-STRENGTH

SEBELA IRELAND LTD	0.025MG; 0.5MG	N017744	001	
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ATROPINE SULFATE; DIPHENOXYLATE HYDROCHLORIDE

CAPSULE; ORAL

DIPHENOXYLATE HYDROCHLORIDE W/ ATROPINE SULFATE

SCHERER RP	0.025MG; 2.5MG	A086440	001	
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SOLUTION; ORAL

COLONALID

MEDPOINTE PHARM HLC	0.025MG/5ML; 2.5MG/5ML	A085735	001	
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LOMANATE

ALPHARMA US PHARMS	0.025MG/5ML; 2.5MG/5ML	A085746	001	
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LOMOTIL

GD SEARLE LLC	0.025MG/5ML; 2.5MG/5ML	N012699	001	
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TABLET; ORAL

COLONALID

MEDPOINTE PHARM HLC	0.025MG; 2.5MG	A085737	001	
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DI-ATRO

MD PHARM	0.025MG; 2.5MG	A085266	001	
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DIPHENOXYLATE HYDROCHLORIDE AND ATROPINE SULFATE

ABLE	0.025MG; 2.5MG	A040395	001	Nov 27, 2000
ASCOT	0.025MG; 2.5MG	A087934	001	Jul 19, 1983
FOSUN PHARMA	0.025MG; 2.5MG	A086173	001	
HEATHER	0.025MG; 2.5MG	A086798	001	
HIKMA PHARMS	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	
PARKE DAVIS	0.025MG; 2.5MG	A087131	001	
PVT FORM	0.025MG; 2.5MG	A085766	001	

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

ATROPINE SULFATE; DIPHENOXYLATE HYDROCHLORIDE

TABLET; ORAL

DIPHENOXYLATE HYDROCHLORIDE AND ATROPINE SULFATE

R AND S PHARMA	0.025MG;2.5MG	A085035	001	
ROXANE	0.025MG;2.5MG	A086057	001	
SUN PHARM INDUSTRIES	0.025MG;2.5MG	A085506	001	
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

MYLAN INSTITUTIONAL	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
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AVOBENZONE; OCTINOXATE; OXYBENZONE

LOTION; TOPICAL

SHADE UVAGUARD

+	BAYER HEALTHCARE LLC	3%;7.5%;3%	N020045	001	Dec 07, 1992
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AZACITIDINE

POWDER; INTRAVENOUS, SUBCUTANEOUS

AZACITIDINE

LUPIN LTD	100MG/VIAL	A210748	001	Feb 27, 2019
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AZATADINE MALEATE

TABLET; ORAL

OPTIMINE

SCHERING	1MG	N017601	001	
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AZATADINE MALEATE; PSEUDOEPHEDRINE SULFATE

TABLET, EXTENDED RELEASE; ORAL

TRINALIN

SCHERING	1MG;120MG	N018506	001	Mar 23, 1982
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AZATHIOPRINE

TABLET; ORAL

IMURAN

+	SEBELA IRELAND LTD	25MG **	N016324	002
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AZATHIOPRINE SODIUM

INJECTABLE; INJECTION

IMURAN

+	CASPER PHARMA LLC	EQ 100MG BASE/VIAL **	N017391	001
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DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

AZELASTINE HYDROCHLORIDE

SOLUTION/DROPS;OPHTHALMIC

OPTIVAR

+ MYLAN SPECIALITY LP 0.05% **

N021127 001 May 22, 2000

SPRAY, METERED;NASAL

ASTEPRO

MYLAN SPECIALITY LP EQ 0.125MG BASE/SPRAY

N022203 001 Oct 15, 2008

AZITHROMYCIN

CAPSULE;ORAL

ZITHROMAX

+ PFIZER EQ 250MG BASE **

N050670 001 Nov 01, 1991

FOR SUSPENSION;ORAL

AZITHROMYCIN

LUPIN LTD EQ 100MG BASE/5ML

A065488 001 May 15, 2015

EQ 200MG BASE/5ML

A065488 002 May 15, 2015

SANDOZ EQ 100MG BASE/5ML

A065297 001 Sep 18, 2006

EQ 200MG BASE/5ML

A065297 002 Sep 18, 2006

FOR SUSPENSION, EXTENDED RELEASE;ORAL

ZMAX

+ PF PRISM CV EQ 2GM BASE/BOT

N050797 001 Jun 10, 2005

INJECTABLE;INJECTION

AZITHROMYCIN

CSPC OUYI EQ 500MG BASE/VIAL

A065265 001 Jan 18, 2007

TEVA PARENTERAL EQ 500MG BASE/VIAL

N050809 001 Dec 19, 2006

EQ 2.5GM BASE/VIAL

N050809 002 Dec 19, 2006

TABLET;ORAL

AZITHROMYCIN

APOTEX CORP EQ 250MG BASE

A065507 001 Jul 13, 2011

EQ 500MG BASE

A065509 001 Jul 13, 2011

EQ 600MG BASE

A065508 001 Jul 13, 2011

MYLAN EQ 250MG BASE

A065365 001 May 30, 2007

EQ 500MG BASE

A065366 001 May 30, 2007

EQ 600MG BASE

A065360 001 Jan 08, 2007

TEVA EQ 250MG BASE

A065153 001 Nov 14, 2005

EQ 600MG BASE

A065150 001 Nov 14, 2005

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

WEST-WARD PHARMS INT 1GM/VIAL

A065286 001 Mar 23, 2011

2GM/VIAL

A065286 002 Mar 23, 2011

DISCONTINUED DRUG PRODUCT LIST

6-50(of 430)

** See List Footnote

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

INJECTABLE;INJECTION

BACITRACIN

MYLAN ASI

50,000 UNITS/VIAL

A090211 001 May 11, 2010

PFIZER

50,000 UNITS/VIAL

A060282 001

PHARMACIA AND UPJOHN

10,000 UNITS/VIAL

A060733 001

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

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

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

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

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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

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			
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	

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

TABLET; ORAL			
BACLOFEN			
MYLAN	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
TABLET, ORALLY DISINTEGRATING; ORAL			
KEMSTRO			
UCB INC	10MG	N021589 001	Oct 30, 2003
	20MG	N021589 002	Oct 30, 2003

BALSALAZIDE DISODIUM

CAPSULE; ORAL			
BALSALAZIDE DISODIUM			
MYLAN	750MG	A077807 001	Dec 28, 2007
TABLET; ORAL			
BALSALAZIDE DISODIUM			
PAR PHARM INC	1.1GM	A206336 001	Sep 08, 2015
GIAZO			
+	VALEANT PHARMS INTL	1.1GM	N022205 001
			Feb 03, 2012

BARIIUM SULFATE

FOR SUSPENSION; ORAL			
E-Z-CAT DRY			
+	BRACCO	40% (9GM/POUCH)	N208036 003
			Jan 03, 2017

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

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

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

ANI PHARMS INC	5MG	A076333	001	Feb 11, 2004
	10MG	A076333	002	Feb 11, 2004
	20MG	A076333	003	Feb 11, 2004
	40MG	A076333	004	Feb 11, 2004
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
HERITAGE PHARMA	5MG	A076267	001	Feb 11, 2004
	10MG	A076267	002	Feb 11, 2004
	20MG	A076267	003	Feb 11, 2004
	40MG	A076267	004	Feb 11, 2004

BENAZEPRIL HYDROCHLORIDE; HYDROCHLOROTHIAZIDE

TABLET; ORAL

BENAZEPRIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE

ANI PHARMS INC	5MG; 6.25MG	A076342	001	Feb 11, 2004
	5MG; 6.25MG	A076348	001	Feb 11, 2004
	10MG; 12.5MG	A076342	002	Feb 11, 2004
	10MG; 12.5MG	A076348	002	Feb 11, 2004
	20MG; 12.5MG	A076342	003	Feb 11, 2004
	20MG; 12.5MG	A076348	003	Feb 11, 2004
	20MG; 25MG	A076342	004	Feb 11, 2004
	20MG; 25MG	A076348	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				
+ US PHARMS HOLDINGS I	5MG; 6.25MG **	N020033	001	May 19, 1992

BENDAMUSTINE HYDROCHLORIDE

SOLUTION; IV (INFUSION)

TREANDA

+ CEPHALON	45MG/0.5ML (90MG/ML)	N022249	003	Sep 13, 2013
+ CEPHALON	180MG/2ML (90MG/ML)	N022249	004	Sep 13, 2013

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

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

NADOLOL AND BENDROFLUMETHIAZIDE

MYLAN 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

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

BENZOYL PEROXIDE; CLINDAMYCIN PHOSPHATE

GEL; TOPICAL

BENZACLIN

BAUSCH 5%; EQ 1% BASE N050756 002 Apr 20, 2007

CLINDAMYCIN PHOSPHATE AND BENZOYL PEROXIDE

ANDA REPOSITORY 2.5%; EQ 1.2% BASE A207194 001 Aug 19, 2019

TARO 3.75%; EQ 1.2% BASE A208683 001 Jun 05, 2018

BENZOYL PEROXIDE; ERYTHROMYCIN

GEL; TOPICAL

AKTIPAK

+ BIOFRONTERA 5%; 3% N050769 001 Nov 27, 2000

BENZPHETAMINE HYDROCHLORIDE

TABLET; ORAL

BENZPHETAMINE HYDROCHLORIDE

EPIC PHARMA LLC 50MG A040714 001 Oct 29, 2007

IMPAX LABS 50MG A040845 001 Nov 18, 2008

TEDOR PHARM 25MG A040747 002 Nov 20, 2015

50MG A040747 001 Mar 30, 2007

DIDREX

+ PHARMACIA AND UPJOHN 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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

BENZTHIAZIDE

TABLET; ORAL

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
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TABLET; ORAL

BENZTROPINE MESYLATE

CHARTWELL RX	1MG	A081265 002	Jan 23, 1992
	2MG	A081265 001	Jan 23, 1992
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
UPSHER SMITH LABS	0.5MG	A040103 001	Dec 12, 1996
	1MG	A040103 002	Dec 12, 1996
	2MG	A040103 003	Dec 12, 1996
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
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BENZYL BENZOATE

EMULSION; TOPICAL

BENZYL BENZOATE

LANNETT	50%	A084535 001	
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BEPOTASTINE BESILATE

SOLUTION/DROPS; OPHTHALMIC

BEPOTASTINE BESILATE

APOTEX	1.5%	A206066 001	Mar 05, 2019
MYLAN	1.5%	A206220 001	Mar 18, 2019

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

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

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
DIPROSONE			
SCHERING	EQ 0.05% BASE		N017536 001
CREAM, AUGMENTED; TOPICAL			
DIPROLENE			
SCHERING	EQ 0.05% BASE		N019408 001 Jan 31, 1986
DIPROLENE AF			
+ MERCK SHARP DOHME	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			
ACP NIMBLE	EQ 0.05% BASE		A071882 001 Jun 06, 1988
ALPHARMA US PHARMS	EQ 0.05% BASE		A071085 001 Feb 03, 1987
PHARMADERM	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			
DIPROLENE			
+ MERCK SHARP DOHME	EQ 0.05% BASE		N019716 001 Aug 01, 1988
OINTMENT; TOPICAL			
ALPHATREX			
SAVAGE LABS	EQ 0.05% BASE		N019143 001 Sep 04, 1984

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

BETAMETHASONE DIPROPIONATE

OINTMENT; TOPICAL

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	

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

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

ACP NIMBLE	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
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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

DISCONTINUED DRUG PRODUCT LIST

6-57(of 430)

** See List Footnote

BETAXOLOL HYDROCHLORIDE; Pilocarpine Hydrochloride

SUSPENSION/DROPS;OPHTHALMIC

BETOPTIC PILO

ALCON

EQ 0.25% BASE;1.75%

N020619 001 Apr 17, 1997

BETAZOLE HYDROCHLORIDE

INJECTABLE;INJECTION

HISTALOG

LILLY

50MG/ML

N009344 001

BETHANECHOL CHLORIDE

INJECTABLE;INJECTION

URECHOLINE

+ ODYSSEY PHARMS

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

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

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

MYOTONACHOL

GLENWOOD

5MG

A084188 001

10MG

A084188 003

25MG

A084188 004

URECHOLINE

ODYSSEY PHARMS

5MG

A089095 001 Dec 19, 1985

+

5MG **

N006536 003

10MG

A088440 001 May 29, 1984

+

10MG **

N006536 002

25MG

A088441 001 May 29, 1984

+

25MG **

N006536 004

+

50MG **

N006536 005

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

BETHANIDINE SULFATE

TABLET; ORAL

TENATHAN

ROBINS AH	10MG	N017675 001
	25MG	N017675 002

BICALUTAMIDE

TABLET; ORAL

BICALUTAMIDE

FRESENIUS KABI USA	50MG	A079045 001	May 13, 2010
KUDCO IRELAND	50MG	A077995 001	Jul 06, 2009
MYLAN	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

HI-TECH PHARMA CO	0.03%	A203299 001	Nov 08, 2018
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LUMIGAN

+ ALLERGAN	0.03% **	N021275 001	Mar 16, 2001
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BIPERIDEN HYDROCHLORIDE

TABLET; ORAL

AKINETON

ABBVIE	2MG	N012003 001
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BIPERIDEN LACTATE

INJECTABLE; INJECTION

AKINETON

ABBVIE	5MG/ML	N012418 002
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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
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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
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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, 5 00MG **	N050719 001	Aug 15, 1996
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BISOPROLOL FUMARATE

TABLET; ORAL

BISOPROLOL FUMARATE

MYLAN	5MG	A075831 001	Dec 14, 2005
	10MG	A075831 002	Dec 14, 2005

ZEBETA

+ TEVA WOMENS	5MG **	N019982 002	Jul 31, 1992
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+	10MG **	N019982 001	Jul 31, 1992
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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
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
MYLAN	2.5MG; 6.25MG	A075768 001	Sep 25, 2000
	5MG; 6.25MG	A075768 002	Sep 25, 2000
	10MG; 6.25MG	A075768 003	Sep 25, 2000
SANDOZ	2.5MG; 6.25MG	A075527 001	Sep 25, 2000
	5MG; 6.25MG	A075527 003	Sep 25, 2000

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

BISOPROLOL FUMARATE; HYDROCHLOROTHIAZIDE

TABLET; ORAL

BISOPROLOL FUMARATE AND HYDROCHLOROTHIAZIDE

	10MG; 6.25MG	A075527	002	Sep 25, 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
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SOLUTION; INHALATION

TORNALATE

SANOFI AVENTIS US	0.2%	N019548	001	Feb 19, 1992
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BIVALIRUDIN

INJECTABLE; INTRAVENOUS

BIVALIRUDIN

APOTEX	250MG/VIAL	A204876	001	Jul 06, 2017
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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

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

BOCEPREVIR

CAPSULE; ORAL

VICTRELIS

MERCK SHARP DOHME	200MG	N202258	001	May 13, 2011
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BORTEZOMIB

POWDER; INTRAVENOUS, SUBCUTANEOUS

BORTEZOMIB

+ HOSPIRA INC	1MG/VIAL	N209191	002	Dec 28, 2018
+	2.5MG/VIAL	N209191	001	Jul 12, 2018

BOSENTAN

TABLET; ORAL

BOSENTAN

ALVOGEN PINE BROOK	62.5MG	A206002	001	Apr 26, 2019
	125MG	A206002	002	Apr 26, 2019
MYLAN	62.5MG	A205173	001	Jan 15, 2020
	125MG	A205173	002	Jan 15, 2020

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
EUROHLTH INTL SARL	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
WEST-WARD PHARMS INT	50MG/ML	A070545	001	May 14, 1986
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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

BRETYLIUM TOSYLATE

INJECTABLE; INJECTION

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	

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

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
COASTAL PHARMS	EQ 0.09% ACID	A201211 001	May 11, 2011
VISTAPHARM	EQ 0.09% ACID	A201941 001	Feb 10, 2015
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
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BROMODIPHENHYDRAMINE HYDROCHLORIDE

CAPSULE; ORAL

AMBODRYL

PARKE DAVIS	25MG	N007984 001	
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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	
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TABLET; ORAL

BROMPHENIRAMINE MALEATE

BARR	4MG	A084468 001	
IVAX SUB TEVA PHARMS	4MG	A084351 001	
NEWTRON PHARMS	4MG	A086987 001	

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

BROMPHENIRAMINE MALEATE

TABLET; ORAL

BROMPHENIRAMINE MALEATE

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	
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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
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BROMFED-DM

WOCKHARDT	2MG/5ML; 10MG/5ML; 30MG/5ML	A089681	001	Dec 22, 1988
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BROMPHENIRAMINE MALEATE, PSEUDOEPHEDRINE HYDROCHLORIDE AND DEXTROMETHORPHAN HYDROBROMIDE

VINTAGE PHARMS	2MG/5ML; 10MG/5ML; 30MG/5ML	A202940	001	Jul 21, 2014
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DIMETANE-DX

+ ROBINS AH	2MG/5ML; 10MG/5ML; 30MG/5ML **	N019279	001	Aug 24, 1984
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BROMPHENIRAMINE MALEATE; PSEUDOEPHEDRINE HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL

EFIDAC 24 PSEUDOEPHEDRINE HYDROCHLORIDE/BROMPHENIRAMINE MALEATE

ALZA	16MG; 240MG	N019672	001	Mar 29, 1996
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BUCLIZINE HYDROCHLORIDE

TABLET; ORAL

BUCLADIN-S

STUART PHARMS	50MG	N010911	006	
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BUDESONIDE

AEROSOL, METERED; NASAL

RHINOCORT

ASTRAZENECA	0.032MG/INH	N020233	001	Feb 14, 1994
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POWDER, METERED; INHALATION

PULMICORT

ASTRAZENECA	0.16MG/INH	N020441	002	Jun 24, 1997
	0.32MG/INH	N020441	003	Jun 24, 1997

BUMETANIDE

INJECTABLE; INJECTION

BUMETANIDE

ATHENEX INC	0.25MG/ML	A074441	001	Jan 27, 1995
HOSPIRA	0.25MG/ML	A074160	001	Oct 30, 1997
TEVA PARENTERAL	0.25MG/ML	A074613	001	Nov 18, 1997

BUMEX

+ VALIDUS PHARMS	0.25MG/ML **	N018226	001	Feb 28, 1983
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BUPIVACAINE HYDROCHLORIDE

INJECTABLE; INJECTION

BUPIVACAINE HYDROCHLORIDE

HOSPIRA	0.75%	A070587	001	Mar 03, 1987
MYLAN ASI	0.25%	A091503	001	Oct 18, 2011
	0.5%	A091503	002	Oct 18, 2011
NOVOCOL HEALTHCARE	0.5%	A211096	001	Feb 19, 2019

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
MYLAN ASI	0.25%	A091487	002	Oct 18, 2011

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

BUPIVACAINE HYDROCHLORIDE

INJECTABLE; INJECTION

BUPIVACAINE HYDROCHLORIDE PRESERVATIVE FREE

0.5%

A091487 001 Oct 18, 2011

0.75%

A091487 003 Oct 18, 2011

INJECTABLE; SPINAL

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

BUPRENORPHINE HYDROCHLORIDE

TABLET; SUBLINGUAL

BUPRENORPHINE HYDROCHLORIDE

BARR

EQ 2MG BASE

A090360 001 May 07, 2010

EQ 8MG BASE

A090360 002 May 07, 2010

MYLAN

EQ 2MG BASE

A201066 001 Mar 06, 2015

EQ 8MG BASE

A201066 002 Mar 06, 2015

SUBUTEX

+ INDIVIOR INC

EQ 2MG BASE **

N020732 002 Oct 08, 2002

+

EQ 8MG BASE **

N020732 003 Oct 08, 2002

BUPRENORPHINE HYDROCHLORIDE; NALOXONE HYDROCHLORIDE

FILM; SUBLINGUAL

CASSIPA

+ TEVA PHARMS USA

EQ 16MG BASE; EQ 4MG BASE

N208042 001 Sep 07, 2018

TABLET; SUBLINGUAL

BUPRENORPHINE HYDROCHLORIDE AND NALOXONE HYDROCHLORIDE

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 INC

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

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

IMPAX LABS

300MG

A077415 002 Dec 15, 2006

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

MYLAN

100MG

A090325 001 Apr 08, 2010

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

BUPROPION HYDROCHLORIDETABLET, EXTENDED RELEASE;ORAL
BUPROPION HYDROCHLORIDE

	150MG	A090325 002	Apr 08, 2010
	150MG	A090941 001	May 03, 2010
	150MG	A090942 001	Jul 14, 2010
	200MG	A090325 003	Apr 08, 2010
	300MG	A090942 002	Jul 14, 2010
SANDOZ	100MG	A076845 001	Jul 14, 2005
	150MG	A076834 001	Jul 14, 2005
	150MG	A076845 002	Jul 14, 2005
SUN PHARM	150MG	A200695 001	Dec 18, 2014
WOCKHARDT LTD	100MG	A201331 001	Aug 30, 2012
	150MG	A201331 002	Aug 30, 2012
	200MG	A201331 003	Aug 30, 2012
YICHANG HUMANWELL	100MG	A211347 001	Oct 16, 2018
	150MG	A211347 002	Oct 16, 2018
	200MG	A211347 003	Oct 16, 2018
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

EGIS	5MG	A075119 001	Mar 14, 2002
	10MG	A075119 002	Mar 14, 2002
	15MG	A075119 003	Jan 23, 2003
FOSUN PHARMA	5MG	A075413 001	Mar 19, 2002
	10MG	A075413 002	Mar 19, 2002
	15MG	A075413 003	Mar 19, 2002
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
	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
OXFORD PHARMS	5MG	A075388 001	May 09, 2002
	10MG	A075388 002	May 09, 2002
	15MG	A075388 003	May 09, 2002
RUBICON	5MG	A075521 001	Apr 05, 2002
	10MG	A075521 002	Apr 05, 2002
	15MG	A075521 003	Apr 05, 2002

BUSULFAN

INJECTABLE; INJECTION

BUSULFAN

MYLAN LABS LTD	6MG/ML	A205184 001	Jul 13, 2018
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DISCONTINUED DRUG PRODUCT LIST

6-64(of 430)

** See List Footnote

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
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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

MYLAN SPECIALITY LP	15MG	N000793 002
	50MG	N000793 003
	100MG	N000793 005

SARISOL NO. 1

HALSEY	15MG	A084719 001
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SARISOL NO. 2

HALSEY	30MG	A084719 002
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SODIUM BUTABARBITAL

HIKMA PHARMS	15MG	A085418 001
	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-TC

MYLAN	1%	N021408 001	Oct 17, 2002
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BUTOCONAZOLE NITRATE

CREAM; VAGINAL

BUTOCONAZOLE NITRATE

PERRIGO PHARMA INTL	2%	N019881 001	Feb 07, 1997
FEMSTAT			
ROCHE PALO	2%	N019215 001	Nov 25, 1985
FEMSTAT 3			
+ BAYER	2%	N020421 001	Dec 21, 1995

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

6-65(of 430)

** See List Footnote

BUTOCONAZOLE NITRATESUPPOSITORY;VAGINAL
FEMSTAT

ROCHE PALO	100MG	N019359	001	Nov 25, 1985
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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
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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
MYLAN	0.5MG	A202947	001	Dec 02, 2013

DOSTINEX

+ PHARMACIA AND UPJOHN 0.5MG **

N020664 001 Dec 23, 1996

CAFFEINE CITRATE

SOLUTION;ORAL

CAFFEINE CITRATE

AM REGENT	EQ 30MG BASE/3ML (EQ 10MG BASE/ML)	A090064	001	Nov 20, 2009
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CAFFEINE; ERGOTAMINE TARTRATE

SUPPOSITORY;RECTAL

CAFERGOT

+ NOVARTIS	100MG;2MG **	N009000	002	
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TABLET;ORAL

CAFERGOT

NOVARTIS	100MG;1MG	N006620	001	
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WIGRAINE

ORGANON USA INC	100MG;1MG	A086562	001	
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CALCIFEDIOL

CAPSULE;ORAL

CALDEROL

ORGANON USA INC	0.02MG	N018312	001	
	0.05MG	N018312	002	

CALCIPOTRIENE

OINTMENT;TOPICAL

DOVONEX

+ LEO PHARMA AS	0.005% **	N020273	001	Dec 29, 1993
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SOLUTION;TOPICAL

DOVONEX

+ LEO PHARM	0.005% **	N020611	001	Mar 03, 1997
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DISCONTINUED DRUG PRODUCT LIST

6-66(of 430)

** See List Footnote

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

+ MYLAN IRELAND LTD

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

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

AKORN

0.002MG/ML

A078066 002 Jan 29, 2008

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

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

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

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; RISEDRONATE SODIUM

TABLET, TABLET; ORAL

ACTONEL WITH CALCIUM (COPACKAGED)

+ WARNER CHILCOTT

EQ 500MG BASE, N/A; N/A, 35MG **

N021823 001 Aug 12, 2005

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

CALCIUM CHLORIDE; DEXTROSE; GLUTATHIONE DISULFIDE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE; SODIUM PHOSPHATE

SOLUTION; IRRIGATION

METHOTREXATE SODIUM

AKORN

0.154MG/ML; 0.92MG/ML; 0.184MG/ML; 0.2MG/ML; 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; 3.05GM/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; 2.21GM/1000ML; 7.07GM/1000ML (5000ML)

N021703 012 Oct 10, 2008

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; OXIGLUTATHIONE; 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

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

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	
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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

DIALYTE CONCENTRATE W/ DEXTROSE 50% IN PLASTIC CONTAINER

B	BRAUN	510MG/100ML;50GM/100ML;200MG/100ML;9.2GM/100ML;9.6GM/100ML	N018807	002	Aug 26, 1983
		510MG/100ML;50GM/100ML;200MG/100ML;9.4GM/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
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DIALYTE W/ DEXTROSE 1.5% IN PLASTIC CONTAINER

B	BRAUN	29MG/100ML;1.5GM/100ML;15MG/100ML;610MG/100ML;560MG/100ML	N018460	001	
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DIALYTE W/ DEXTROSE 4.25% IN PLASTIC CONTAINER

B	BRAUN	29MG/100ML;4.25GM/100ML;15MG/100ML;610MG/100ML;560MG/100ML	N018460	003	
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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;567MG/100ML;392MG/100ML	N018379	002	
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DELFLX W/ DEXTROSE 2.5% IN PLASTIC CONTAINER

	FRESENIUS MEDCL	25.7MG/100ML;2.5GM/100ML;15.2MG/100ML;567MG/100ML;392MG/100ML	N018379	003	
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DELFLX W/ DEXTROSE 3.5% IN PLASTIC CONTAINER

	FRESENIUS MEDCL	25.7MG/100ML;3.5GM/100ML;15.2MG/100ML;567MG/100ML;392MG/100ML	N018379	007	Jun 24, 1988
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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	
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DELFLX-LM W/ DEXTROSE 1.5% IN PLASTIC CONTAINER

	FRESENIUS MEDCL	25.7MG/100ML;1.5GM/100ML;5.08MG/100ML;538MG/100ML;448MG/100ML	N018379	004	Jul 07, 1982
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DELFLX-LM W/ DEXTROSE 2.5% IN PLASTIC CONTAINER

	FRESENIUS MEDCL	25.7MG/100ML;2.5GM/100ML;5.08MG/100ML;538MG/100ML;448MG/100ML	N018379	005	Jul 07, 1982
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DELFLX-LM W/ DEXTROSE 3.5% IN PLASTIC CONTAINER

	FRESENIUS MEDCL	25.7MG/100ML;3.5GM/100ML;5.08MG/100ML;538MG/100ML;448MG/100ML	N018379	008	Jun 24, 1988
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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
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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
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		26MG/100ML;1.5GM/100ML;15MG/100ML;560MG/100ML;390MG/100ML	N018460	002	
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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
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		26MG/100ML;5GM/100ML;5MG/100ML;530MG/100ML;450MG/100ML	N018460	008	Jan 29, 1986
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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
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		26MG/100ML;4.25GM/100ML;15MG/100ML;560MG/100ML;390MG/100ML	N018460	004	
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DIANEAL 137 W/ DEXTROSE 1.5% IN PLASTIC CONTAINER

	BAXTER HLTHCARE	25.7MG/100ML;1.5GM/100ML;15.2MG/100ML;567MG/100ML;392MG/100ML	N017512	001	
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Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

CALCIUM CHLORIDE; DEXTROSE; MAGNESIUM CHLORIDE; SODIUM CHLORIDE; SODIUM LACTATE

SOLUTION; INTRAPERITONEAL

DIANEAL 137 W/ DEXTROSE 2.5% IN PLASTIC CONTAINER				
BAXTER HLTHCARE	25.7MG/100ML;2.5GM/100ML;15.2MG/100ML;5	N017512	003	
	67MG/100ML;392MG/100ML			
DIANEAL 137 W/ DEXTROSE 4.25% IN PLASTIC CONTAINER				
BAXTER HLTHCARE	25.7MG/100ML;4.25GM/100ML;15.2MG/100ML;	N017512	002	
	567MG/100ML;392MG/100ML			
DIANEAL LOW CALCIUM W/ DEXTROSE 3.5% IN PLASTIC CONTAINER				
+ BAXTER HLTHCARE	18.3MG/100ML;3.5GM/100ML;5.08MG/100ML;5	N020183	003	Dec 04, 1992
	38MG/100ML;448MG/100ML			
DIANEAL PD-1 W/ DEXTROSE 1.5% IN PLASTIC CONTAINER				
BAXTER HLTHCARE	25.7MG/100ML;1.5GM/100ML;15.2MG/100ML;5	N017512	007	Jul 09, 1984
	67MG/100ML;392MG/100ML			
DIANEAL PD-1 W/ DEXTROSE 2.5% IN PLASTIC CONTAINER				
BAXTER HLTHCARE	25.7MG/100ML;2.5GM/100ML;15.2MG/100ML;5	N017512	008	Jul 09, 1984
	67MG/100ML;392MG/100ML			
DIANEAL PD-1 W/ DEXTROSE 3.5% IN PLASTIC CONTAINER				
BAXTER HLTHCARE	25.7MG/100ML;3.5GM/100ML;15.2MG/100ML;5	N017512	010	Nov 18, 1985
	67MG/100ML;392MG/100ML			
DIANEAL PD-1 W/ DEXTROSE 4.25% IN PLASTIC CONTAINER				
BAXTER HLTHCARE	25.7MG/100ML;4.25GM/100ML;15.2MG/100ML;	N017512	009	Jul 09, 1984
	567MG/100ML;392MG/100ML			
DIANEAL PD-2 W/ DEXTROSE 3.5% IN PLASTIC CONTAINER				
+ BAXTER HLTHCARE	25.7MG/100ML;3.5GM/100ML;5.08MG/100ML;5	N017512	011	Nov 18, 1985
	38MG/100ML;448MG/100ML			
INPERSOL-LC/LM W/ DEXTROSE 1.5% IN PLASTIC CONTAINER				
FRESENIUS	18.4MG/100ML;1.5GM/100ML;5.08MG/100ML;5	A020374	001	Jun 13, 1994
	38MG/100ML;448MG/100ML			
INPERSOL-LC/LM W/ DEXTROSE 2.5% IN PLASTIC CONTAINER				
FRESENIUS	18.4MG/100ML;2.5GM/100ML;5.08MG/100ML;5	A020374	002	Jun 13, 1994
	38MG/100ML;448MG/100ML			
INPERSOL-LC/LM W/ DEXTROSE 3.5% IN PLASTIC CONTAINER				
FRESENIUS	18.4MG/100ML;3.5GM/100ML;5.08MG/100ML;5	A020374	003	Jun 13, 1994
	38MG/100ML;448MG/100ML			
INPERSOL-LC/LM W/ DEXTROSE 4.25% IN PLASTIC CONTAINER				
FRESENIUS	18.4MG/100ML;4.25GM/100ML;5.08MG/100ML;	A020374	004	Jun 13, 1994
	538MG/100ML;448MG/100ML			

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/1	N018258	001	
	00ML;600MG/100ML			

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/1	N018254	001	
	00ML			
DEXTROSE 5% IN RINGER'S IN PLASTIC CONTAINER				
B BRAUN	33MG/100ML;5GM/100ML;30MG/100ML;860MG/1	N018256	001	
	00ML			
	33MG/100ML;5GM/100ML;30MG/100ML;860MG/1	N020000	001	Apr 17, 1992
	00ML			
BAXTER HLTHCARE	33MG/100ML;5GM/100ML;30MG/100ML;860MG/1	N016695	001	
	00ML			

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/100	N019634	002	Feb 24, 1988
	ML;62MG/100ML			
DEXTROSE 5% IN LACTATED RINGER'S IN PLASTIC CONTAINER				
B BRAUN	20MG/100ML;5GM/100ML;30MG/100ML;600MG/1	N017510	001	
	00ML;310MG/100ML			
MILES	20MG/100ML;5GM/100ML;30MG/100ML;600MG/1	N018499	001	
	00ML;310MG/100ML			
POTASSIUM CHLORIDE 10MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER				
+ ICU MEDICAL INC	20MG/100ML;5GM/100ML;104MG/100ML;600MG/	N019685	005	Oct 17, 1988
	100ML;310MG/100ML			
+ ICU MEDICAL INC	20MG/100ML;5GM/100ML;179MG/100ML;600MG/	N019685	006	Oct 17, 1988
	100ML;310MG/100ML			
POTASSIUM CHLORIDE 15MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER				
+ ICU MEDICAL INC	20MG/100ML;5GM/100ML;254MG/100ML;	N019685	007	Oct 17, 1988

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

CALCIUM CHLORIDE; DEXTROSE; POTASSIUM CHLORIDE; SODIUM CHLORIDE; SODIUM LACTATE

INJECTABLE; INJECTION

POTASSIUM CHLORIDE 15MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER			
	600MG/100ML; 310MG/100ML		
POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER			
+ 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
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

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	

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

CALCIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM CHLORIDE; SODIUM LACTATE

SOLUTION; IRRIGATION

LACTATED RINGER'S IN PLASTIC CONTAINER

BAXTER HLTHCARE	20MG/100ML; 30MG/100ML; 600MG/100ML; 310MG/100ML	N019933 001	Aug 29, 1989
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CALCIUM GLUCEPTATE

INJECTABLE; INJECTION

CALCIUM GLUCEPTATE

ABBOT	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	
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CALCIUM; MEGLUMINE; METRIZOIC ACID

INJECTABLE; INJECTION

ISOPAQUE 280

GE HEALTHCARE	0.35MG/ML; 140.1MG/ML; 461.8MG/ML	N017506 001	
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CANDESARTAN CILEXETIL

TABLET; ORAL

CANDESARTAN CILEXETIL

APOTEX INC	4MG	A202079 001	Jan 10, 2014
	8MG	A202079 002	Jan 10, 2014
	16MG	A202079 003	Jan 10, 2014
	32MG	A202079 004	Jan 10, 2014

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

VANOVID

SANOFI AVENTIS US	0.6MG/GM	A061596 001	
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TABLET; VAGINAL

VANOVID

SANOFI AVENTIS US	3MG	A061613 001	
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CAPTOPRIL

TABLET; ORAL

CAPOTEN

+	PAR PHARM	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

CAPTOPRIL

ACP NIMBLE	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
APOTHECON	12.5MG	A074472 001	Mar 31, 1995
	25MG	A074472 002	Mar 31, 1995
	50MG	A074472 003	Mar 31, 1995

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

CAPTOPRILTABLET; ORAL
CAPTOPRIL

	100MG	A074472	004	Mar 31, 1995
COREPHARMA	12.5MG	A074737	001	Oct 28, 1998
	25MG	A074737	002	Oct 28, 1998
	50MG	A074737	003	Oct 28, 1998
	100MG	A074737	004	Oct 28, 1998
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
PAR PHARM	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
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
WATSON LABS	12.5MG	A074451	001	Feb 13, 1996
	12.5MG	A074576	001	Apr 23, 1996
	25MG	A074451	002	Feb 13, 1996
	25MG	A074576	002	Apr 23, 1996
	50MG	A074451	003	Feb 13, 1996
	50MG	A074576	003	Apr 23, 1996
	100MG	A074451	004	Feb 13, 1996
	100MG	A074576	004	Apr 23, 1996
YAOPHARMA CO LTD	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

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				
ACP NIMBLE	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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

CAPTOPRIL; HYDROCHLOROTHIAZIDE

TABLET; ORAL

CAPTOPRIL AND HYDROCHLOROTHIAZIDE

	50MG;15MG	A075055 004	Jun 18, 1998
	50MG;25MG	A075055 003	Jun 18, 1998
VINTAGE PHARMS LLC	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

CAPSULE, EXTENDED RELEASE; ORAL

CARBAMAZEPINE

NOSTRUM LABS INC	100MG	A076697 001	May 20, 2011
	200MG	A076697 002	May 20, 2011
	300MG	A076697 003	May 20, 2011

SOLUTION; INTRAVENOUS

CARNEXIV

+ LUNDBECK PHARMS LLC	200MG/20ML (10MG/ML)	N206030 001	Oct 07, 2016
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SUSPENSION; ORAL

CARBAMAZEPINE

TARO	100MG/5ML	A075875 001	Dec 21, 2000
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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
USL PHARMA	200MG	A070300 001	May 15, 1986
WARNER CHILCOTT	200MG	A070429 001	Jan 02, 1987

TERIL

TARO	200MG	A076525 001	Sep 26, 2003
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TABLET, CHEWABLE; ORAL

CARBAMAZEPINE

JUBILANT CADISTA	100MG	A071940 001	Feb 01, 1988
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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	
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CARBIDOPA; ENTACAPONE; LEVODOPA

TABLET; ORAL

CARBIDOPA, LEVODOPA AND ENTACAPONE

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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

CARBIDOPA; LEVODOPA

TABLET; ORAL

CARBIDOPA AND LEVODOPA

ANI PHARMS INC	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

TABLET, EXTENDED RELEASE; ORAL

CARBIDOPA AND LEVODOPA

KV PHARM	50MG;200MG	A076663 001	Jun 24, 2004
SINEMET CR			
+ MERCK SHARP DOHME	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
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	
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SOLUTION; ORAL

CARBINOXAMINE MALEATE

CYPRESS PHARM	4MG/5ML	A090418 001	May 04, 2010
VINTAGE PHARMS	4MG/5ML	A040814 001	Feb 26, 2008

TABLET; ORAL

CARBINOXAMINE MALEATE

CYPRESS PHARM	4MG	A090417 001	Aug 23, 2010
VINTAGE PHARMS	4MG	A040639 002	May 30, 2008

CLISTIN

+ ORTHO MCNEIL PHARM	4MG **	N008915 001	
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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
HOSPIRA	50MG/VIAL	A076473 001	Oct 27, 2004
	150MG/VIAL	A076473 002	Oct 27, 2004
	450MG/VIAL	A076473 003	Oct 27, 2004
MYLAN LABS LTD	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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

6-75(of 430)

** See List Footnote

CARBOPLATIN

INJECTABLE; INJECTION

CARBOPLATIN

	150MG/VIAL	A076162 002	Oct 14, 2004
	450MG/VIAL	A076162 003	Oct 14, 2004
WEST-WARD PHARMS INT	50MG/VIAL	A076099 001	Oct 14, 2004
	150MG/VIAL	A076099 002	Oct 14, 2004
	450MG/VIAL	A076099 003	Oct 14, 2004
PARAPLATIN			
+	CORDEN PHARMA	50MG/VIAL **	N019880 001
+		150MG/VIAL **	N019880 002
+		450MG/VIAL **	N019880 003

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
MYLAN INSTITUTIONAL			
	50MG/5ML (10MG/ML)	A077998 001	Apr 24, 2007
	150MG/15ML (10MG/ML)	A077998 002	Apr 24, 2007
	450MG/45ML (10MG/ML)	A077998 003	Apr 24, 2007
MYLAN LABS LTD			
	50MG/5ML (10MG/ML)	A091063 001	Nov 09, 2011
	150MG/15ML (10MG/ML)	A091063 002	Nov 09, 2011
	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
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
PARAPLATIN			
+	CORDENPHARMA	50MG/5ML (10MG/ML) **	N020452 001
+		150MG/15ML (10MG/ML) **	N020452 002
+		450MG/45ML (10MG/ML) **	N020452 003
+		600MG/60ML (10MG/ML) **	N020452 004

CARISOPRODOL

CAPSULE; ORAL

SOMA

+	MYLAN SPECIALITY LP	250MG	N011792 003
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TABLET; ORAL

CARISOPRODOL

ABLE			
	350MG	A040421 001	Jun 21, 2001
EPIC PHARMA LLC			
	350MG	A040397 001	Sep 21, 2000
FOSUN PHARMA			
	350MG	A081025 001	Apr 13, 1989
HIKMA INTL PHARMS			
	350MG	A040124 001	Jan 24, 1996
OXFORD PHARMS			
	350MG	A040188 001	Mar 07, 1997
PIONEER PHARMS			
	350MG	A089390 001	Oct 13, 1988
SANDOZ			
	350MG	A089566 001	Aug 30, 1988
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	

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

CARPHENAZINE 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

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

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

WOCKHARDT LTD

3.125MG

A078786 001 Dec 22, 2009

6.25MG

A078786 002 Dec 22, 2009

12.5MG

A078786 003 Dec 22, 2009

25MG

A078786 004 Dec 22, 2009

CASPOFUNGIN ACETATE

POWDER; INTRAVENOUS

CASPOFUNGIN ACETATE

CIPLA

50MG/VIAL

A209489 001 Jul 12, 2018

70MG/VIAL

A209489 002 Jul 12, 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

DAVA PHARMS INC

EQ 250MG BASE

A064107 001 Apr 27, 1995

EQ 500MG BASE

A064107 002 Apr 27, 1995

HERITAGE PHARMA

EQ 250MG BASE

A064148 001 May 23, 1996

EQ 500MG BASE

A064148 002 May 23, 1996

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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

CEFACLOR

CAPSULE; ORAL

CEFACLOR

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

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

CEFACLOR

WORLD GEN

EQ 500MG BASE

A065057 001 Jan 05, 2001

CEFADROXIL/CEFADROXIL HEMIHYDRATE

CAPSULE; ORAL

CEFADROXIL

CSPC OUYI

EQ 500MG BASE

A205072 001 Jul 28, 2017

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 INC

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

SUN PHARM INDS LTD

EQ 125MG BASE/5ML

A065115 001 Mar 26, 2003

EQ 250MG BASE/5ML

A065115 002 Mar 26, 2003

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

CEFADROXIL/CEFADROXIL HEMIHYDRATE

FOR SUSPENSION; ORAL

CEFADROXIL

	EQ 500MG BASE/5ML	A065115 003	Mar 26, 2003
DURICEF			
+ WARNER CHILCOTT	EQ 125MG BASE/5ML **	N050527 002	
+	EQ 250MG BASE/5ML **	N050527 003	
+	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

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
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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

CEFAZOLIN SODIUMINJECTABLE; INJECTION
CEFAZOLIN SODIUM

	EQ 20GM BASE/VIAL	A062989 003	Dec 29, 1989
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
HOSPIRA INC	EQ 10GM BASE/VIAL	A065247 001	Aug 12, 2005
	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 1GM BASE/VIAL	A201654 001	Feb 03, 2016
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
WEST-WARD PHARMS INT	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
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			
FOSUN PHARMA	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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

CEFIXIME

FOR SUSPENSION; ORAL

CEFIXIME

SANDOZ INC

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

CEFMENOXIME

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

CEFOPERAZONE

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

CEFOPERAZONE 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

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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

CEFOTAXIME SODIUM

INJECTABLE; INJECTION

CEFOTAXIME

	EQ 20GM BASE/VIAL	A064201 002	Mar 24, 2000
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 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 1GM BASE/VIAL	A203132 001	Feb 19, 2016
	EQ 2GM BASE/VIAL	A065290 003	Aug 11, 2006
	EQ 2GM BASE/VIAL	A065293 002	Aug 10, 2006
	EQ 2GM BASE/VIAL	A203132 002	Feb 19, 2016
	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
+ US PHARM HOLDINGS	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			
US PHARM HOLDINGS	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			
US PHARM HOLDINGS	EQ 20MG BASE/ML	N050596 001	May 20, 1985
	EQ 40MG BASE/ML	N050596 003	May 20, 1985

CEFOTETAN DISODIUM

INJECTABLE; INJECTION

CEFOTAN

TELIGENT	EQ 10GM BASE/VIAL	N050588 003	Apr 25, 1988
TELIGENT PHARMA INC	EQ 1GM BASE/VIAL	A063293 001	Apr 29, 1993
	EQ 2GM BASE/VIAL	A063293 002	Apr 29, 1993
CEFOTAN IN PLASTIC CONTAINER			
TELIGENT	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
WEST-WARD PHARM CORP	EQ 10GM BASE/VIAL	A091030 001	Oct 26, 2011

CEFOTIAM HYDROCHLORIDE

INJECTABLE; INJECTION

CERADON

TAKEDA	EQ 1GM BASE/VIAL	N050601 001	Dec 30, 1988
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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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

CEFOXITIN SODIUM

INJECTABLE; INJECTION

CEFOXITIN

EQ 10GM BASE/VIAL

A065312 001 Feb 13, 2006

MEFOXIN

MYLAN INSTITUTIONAL

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 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

SANDOZ

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

RANBAXY LABS LTD

125MG/5ML

A065202 001 Jun 30, 2006

250MG/5ML

A065202 002 Jun 30, 2006

CEFZIL

+ CORDEN PHARMA

125MG/5ML **

N050665 001 Dec 23, 1991

+

250MG/5ML **

N050665 002 Dec 23, 1991

TABLET; ORAL

CEFPROZIL

RANBAXY LABS LTD

250MG

A065198 001 Dec 13, 2006

500MG

A065198 002 Dec 13, 2006

CEFZIL

+ CORDEN PHARMA

250MG **

N050664 001 Dec 23, 1991

+

500MG **

N050664 002 Dec 23, 1991

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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

CEFTAZIDIME

INJECTABLE; INJECTION

CEFTAZIDIME

6GM/VIAL

A065482 001 May 28, 2010

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

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

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

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

TELIGENT

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

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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

CEFTRIAXONE SODIUM

INJECTABLE; INJECTION

CEFTRIAXONE

	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			
+	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

INJECTABLE; INTRAMUSCULAR, INTRAVENOUS

CEFTRIAXONE

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
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
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			
+	EQ 250MG BASE/VIAL **	N050585 001	Dec 21, 1984
+	EQ 500MG BASE/VIAL **	N050585 002	Dec 21, 1984
+	EQ 1GM BASE/VIAL **	N050585 003	Dec 21, 1984
+	EQ 2GM BASE/VIAL **	N050585 004	Dec 21, 1984

CEFTRIAXONE SODIUM; LIDOCAINE

INJECTABLE; INJECTION

ROCEPHIN KIT

HOFFMANN LA ROCHE	EQ 500MG BASE/VIAL, N/A; N/A, 1%	N050585 007	May 08, 1996
	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
+		EQ 250MG BASE/5ML	N050672 002	Apr 29, 1997
CEFUROXIME AXETIL				
SUN PHARM INDS LTD	EQ 125MG BASE/5ML	A065323 001	Feb 05, 2008	

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

CEFUROXIME AXETILFOR SUSPENSION; ORAL
CEFUROXIME AXETIL

	EQ 250MG BASE/5ML	A065323 002	Feb 05, 2008
TABLET; ORAL			
CEFTIN			
+	GLAXOSMITHKLINE	EQ 125MG BASE **	N050605 001 Dec 28, 1987
+		EQ 250MG BASE **	N050605 002 Dec 28, 1987
+		EQ 500MG BASE **	N050605 003 Dec 28, 1987
CEFUROXIME AXETIL			
	ANI PHARMS INC	EQ 250MG BASE	A065190 001 Oct 18, 2004
		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 SODIUMINJECTABLE; INJECTION
CEFUROXIME SODIUM

	FRESENIUS KABI USA	EQ 1.5GM BASE/VIAL	A065001 002 May 30, 2001
		EQ 7.5GM BASE/VIAL	A065002 001 Sep 28, 1998
	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 IN PLASTIC CONTAINER			
	TELIGENT	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
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

CELLULOSE SODIUM PHOSPHATEPOWDER; ORAL
CALCIBIND

	MISSION PHARMA	2.5GM/PACKET	N018757 002 Dec 28, 1982
		300GM/BOT	N018757 003 Oct 16, 1984

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

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	
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 333MG BASE **	N050405 004	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
	EQ 250MG BASE/5ML	A062777 001	Aug 06, 1987
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	
+ PRAGMA	EQ 125MG BASE/5ML **	N050406 001	
+ PRAGMA	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	

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

CEPHALEXINTABLET, 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 HYDROCHLORIDETABLET;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

CEPHALOGLYCINCAPSULE;ORAL
KAFOCIN

LILLY	250MG	N050219 001	
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CEPHALOTHIN SODIUMINJECTABLE;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

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 SODIUMINJECTABLE;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	

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

CEPHAPIRIN SODIUM

INJECTABLE; INJECTION

CEFADYL

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
WEST-WARD PHARMS INT 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	

VELOSEF '250'

ERSANA

250MG	N050548	001	
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VELOSEF '500'

ERSANA

500MG	N050548	002	
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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	
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VELOSEF '250'

APOTHECON

250MG/5ML	A061763	002	
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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		
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CERITINIB

CAPSULE; ORAL

ZYKADIA

+ NOVARTIS

150MG

N205755	001	Apr 29, 2014	
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DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

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	
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CETIRIZINE HYDROCHLORIDE

SYRUP; ORAL

CETIRIZINE HYDROCHLORIDE

ACTAVIS MID ATLANTIC	5MG/5ML	A078617 001	Feb 02, 2010
AUROBINDO PHARMA LTD	5MG/5ML	A090751 001	Dec 16, 2009
LANNETT CO INC	5MG/5ML	A078496 001	Sep 25, 2009
PHARM ASSOC	5MG/5ML	A078412 001	Jun 18, 2008
RANBAXY LABS LTD	5MG/5ML	A077472 001	Jun 18, 2008
TORRENT	5MG/5ML	A078870 001	Apr 27, 2009
WOCKHARDT	5MG/5ML	A078757 001	Aug 28, 2009

CHILDREN'S CETIRIZINE HYDROCHLORIDE ALLERGY

ACTAVIS MID ATLANTIC	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
TORRENT	5MG/5ML	A090474 002	Mar 30, 2009

CHILDREN'S CETIRIZINE HYDROCHLORIDE HIVES RELIEF

ACTAVIS MID ATLANTIC	5MG/5ML	A090378 001	May 09, 2008
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
TORRENT	5MG/5ML	A090474 001	Mar 30, 2009

ZYRTEC

J AND J CONSUMER INC	5MG/5ML **	N020346 001	Sep 27, 1996
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TABLET; ORAL

CETIRIZINE HYDROCHLORIDE ALLERGY

HERITAGE PHARMA	5MG	A078615 003	Dec 28, 2007
	10MG	A078615 004	Dec 28, 2007
SUN PHARM INDS INC	5MG	A077499 001	Dec 27, 2007
	10MG	A077499 002	Dec 27, 2007

CETIRIZINE HYDROCHLORIDE HIVES

SUN PHARM INDS INC	5MG	A077499 003	Dec 27, 2007
	10MG	A077499 004	Dec 27, 2007

ZYRTEC ALLERGY

+ J AND J CONSUMER INC	5MG	N019835 003	Nov 16, 2007
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ZYRTEC HIVES RELIEF

+ J AND J CONSUMER INC	5MG	N019835 005	Nov 16, 2007
+ J AND J CONSUMER INC	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 ZYRTEC ALLERGY

+ J AND J CONSUMER INC	5MG **	N021621 003	Nov 16, 2007
+ J AND J CONSUMER INC	10MG **	N021621 004	Nov 16, 2007

CHILDREN'S ZYRTEC HIVES RELIEF

+ J AND J CONSUMER INC	5MG **	N021621 005	Nov 16, 2007
+ J AND J CONSUMER INC	10MG **	N021621 006	Nov 16, 2007

DISCONTINUED DRUG PRODUCT LIST

6-90(of 430)

** See List Footnote

CETIRIZINE HYDROCHLORIDE; PSEUDOEPHEDRINE HYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL

CETIRIZINE HYDROCHLORIDE AND PSEUDOEPHEDRINE HYDROCHLORIDE
PERRIGO R AND D 5MG;120MG

A210719 001 Nov 16, 2018

CETRORELIX

INJECTABLE;INJECTION

CETROTIDE

EMD SERONO INC EQ 3MG BASE/ML

N021197 002 Aug 11, 2000

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

AKORN 0.5%

A062042 001

ALCON 0.5%

A062628 001 Sep 25, 1985

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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CHLORAMPHENICOL SODIUM SUCCINATE

INJECTABLE; INJECTION

CHLORAMPHENICOL

ELKINS SINN EQ 1GM BASE/VIAL

A062406 001 Nov 09, 1982

CHLORAMPHENICOL SODIUM SUCCINATE

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

CHLORAMPHENICOL; DESOXYRIBONUCLEASE; FIBRINOLYSIN

OINTMENT; TOPICAL

ELASE-CHLOROMYCETIN

+ PARKE DAVIS 10MG/GM; 666 UNITS/GM; 1 UNITS/GM

N050294 001

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

CHLOROMYCIN

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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CHLORDIAZEPOXIDE HYDROCHLORIDE

CAPSULE; ORAL

CHLORDIAZEPOXIDE HYDROCHLORIDE

	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	
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	A084678	001	
	5MG	A084919	001	
	10MG	A084041	001	
	10MG	A084920	001	
	25MG	A084679	002	
	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; ESTROGENS, ESTERIFIED

TABLET; ORAL

MENRIUM 10-4

ROCHE 10MG; 0.4MG N014740 006

MENRIUM 5-2

ROCHE 5MG; 0.2MG N014740 002

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

CHLORDIAZEPOXIDE; ESTROGENS, ESTERIFIED

TABLET; ORAL

MENRIUM 5-4

ROCHE

5MG;0.4MG

N014740 004

CHLORHEXIDINE GLUCONATE

SOLUTION; DENTAL

CHLORHEXIDINE GLUCONATE

BAJAJ

0.12%

A075561 001 Nov 14, 2000

SOLUTION; TOPICAL

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

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

NESACAINE-MPF

FRESENIUS KABI USA

2%

N009435 003

3%

N009435 004

CHLOROQUINE HYDROCHLORIDE

INJECTABLE; INJECTION

ARALEN HYDROCHLORIDE

SANOFI AVENTIS US

EQ 40MG BASE/ML

N006002 002

CHLOROQUINE PHOSPHATE

TABLET; ORAL

ARALEN

+ SANOFI AVENTIS US

EQ 300MG BASE

N006002 001

CHLOROQUINE PHOSPHATE

HIKMA PHARMS

EQ 150MG BASE

A083082 001

IMPAX LABS

EQ 150MG BASE

A080880 001

EQ 300MG BASE

A040516 001 Aug 29, 2003

MD PHARM

EQ 150MG BASE

A087228 001

PUREPAC PHARM

EQ 150MG BASE

A080886 001

TEVA

EQ 150MG BASE

A087504 001 Jan 13, 1982

WATSON LABS

EQ 150MG BASE

A087979 001 Dec 21, 1982

EQ 300MG BASE

A088030 001 Dec 21, 1982

CHLOROQUINE PHOSPHATE; PRIMAQUINE PHOSPHATE

TABLET; ORAL

ARALEN PHOSPHATE W/ PRIMAQUINE PHOSPHATE

SANOFI AVENTIS US

EQ 300MG BASE;EQ 45MG BASE

N014860 002

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

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	
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				
+ OAK PHARMS AKORN	250MG **	N011145	004	
+	500MG **	N011145	002	

CHLOROTHIAZIDE; METHYLDOPA

TABLET; ORAL

ALDOCLOR-150

MERCK	150MG;250MG	N016016	001	
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ALDOCLOR-250

MERCK	250MG;250MG	N016016	002	
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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 PHARMS	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

MYLAN	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
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DIUPRES-500

MERCK	500MG;0.125MG	N011635	006	Aug 26, 1987
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CHLOROTRIANISENE

CAPSULE; ORAL

CHLOROTRIANISENE

BANNER PHARMACAPS	12MG	A084652	001	
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TACE

SANOFI AVENTIS US	12MG	N008102	004	
	25MG	N011444	001	
	72MG	N016235	001	

CHLOROXYLINE

SHAMPOO; TOPICAL

CAPITROL

WESTWOOD SQUIBB	2%	N017594	001	
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CHLORPHENESIN CARBAMATE

TABLET; ORAL

MAOLATE

PAMLAB LLC	400MG	N014217	002	
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CHLORPHENIRAMINE MALEATE

CAPSULE, EXTENDED RELEASE; ORAL

CHLORPHENIRAMINE MALEATE

AUROLIFE PHARMA LLC	12MG	A070797	001	Aug 12, 1988
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TELDRIN

GLAXOSMITHKLINE	8MG	N017369	001	
	12MG	N017369	002	

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

CHLORPHENIRAMINE MALEATE

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
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SYRUP; ORAL

CHLOR-TRIMETON

SCHERING	2MG/5ML	N006921	006
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CHLORPHENIRAMINE MALEATE

PHARM ASSOC	2MG/5ML	A087520	001 Feb 10, 1982
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TABLET; ORAL

ANTAGONATE

BAYER PHARMS	4MG	A083381	001
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CHLOR-TRIMETON

SCHERING	4MG	N006921	002
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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
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PHENETRON

LANNETT	4MG	A080846	001
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TABLET, EXTENDED RELEASE; ORAL

CHLOR-TRIMETON

+ BAYER HEALTHCARE LLC	8MG **	N007638	001
+	12MG **	N007638	002

EFIDAC 24 CHLORPHENIRAMINE MALEATE

ALZA	16MG	N019746	002 Nov 18, 1994
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CHLORPHENIRAMINE MALEATE; HYDROCODONE BITARTRATE

SOLUTION; ORAL

HYDROCODONE BITARTRATE AND CHLORPHENIRAMINE MALEATE

ACELLA	4MG/5ML; 5MG/5ML	A206891	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
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DISCONTINUED DRUG PRODUCT LIST

6-96(of 430)

** See List Footnote

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
TORRENT	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
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CHLORPHENIRAMINE MALEATE; PHENYLPROPANOLAMINE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE;ORAL

COLD CAPSULE IV

GRAHAM DM	12MG;75MG	N018793 001	Apr 25, 1985
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COLD CAPSULE V

GRAHAM DM	8MG;75MG	N018794 001	Apr 23, 1985
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TABLET, EXTENDED RELEASE;ORAL

TRIAMINIC-12

NOVARTIS	12MG;75MG	N018115 001	
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CHLORPHENIRAMINE MALEATE; PSEUDOEPHEDRINE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE;ORAL

CODIMAL-L.A. 12

SCHWARZ PHARMA	12MG;120MG	N018935 001	Apr 15, 1985
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ISOCOLOR

FISONS	8MG;120MG	N018747 001	Mar 06, 1986
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PSEUDOEPHEDRINE HYDROCHLORIDE AND CHLORPHENIRAMINE MALEATE

CENT PHARMS	8MG;120MG	N019428 001	Aug 02, 1988
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GRAHAM DM	8MG;120MG	N018844 001	Mar 20, 1985
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	12MG;120MG	N018843 001	Mar 18, 1985
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KV PHARM	12MG;120MG	A071455 001	Mar 01, 1989
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CHLORPHENIRAMINE MALEATE; PSEUDOEPHEDRINE SULFATE

TABLET, EXTENDED RELEASE;ORAL

CHLOR-TRIMETON

+ BAYER HEALTHCARE LLC	8MG;120MG	N018397 001	
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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
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PENNTUSS

FISONS	EQ 4MG MALEATE/5ML;EQ 10MG BASE/5ML	N018928 001	Aug 14, 1985
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CHLORPHENIRAMINE POLISTIREX; HYDROCODONE POLISTIREX

SUSPENSION, EXTENDED RELEASE;ORAL

TUSSIONEX PENNKINETIC

+ UCB INC	EQ 8MG MALEATE/5ML;EQ 10MG BITARTRATE/5ML **	N019111 001	Dec 31, 1987
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CHLORPHENTERMINE HYDROCHLORIDE

TABLET;ORAL

PRE-SATE

PARKE DAVIS	EQ 65MG BASE	N014696 001	
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CHLORPROMAZINE

SUPPOSITORY;RECTAL

THORAZINE

+ GLAXOSMITHKLINE	25MG **	N009149 024	
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+	100MG **	N009149 033	
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CHLORPROMAZINE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE;ORAL

THORAZINE

GLAXOSMITHKLINE	30MG	N011120 016	
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	75MG	N011120 017	
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	150MG	N011120 018	
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	200MG	N011120 019	
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	300MG	N011120 020	
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CONCENTRATE;ORAL

CHLORPROMAZINE HYDROCHLORIDE

ACTAVIS MID ATLANTIC	100MG/ML	A086863 001	
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PHARM ASSOC	30MG/ML	A040231 001	Dec 30, 1999
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	100MG/ML	A040224 001	Jan 26, 1999
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Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

CHLORPROMAZINE HYDROCHLORIDE

CONCENTRATE; ORAL

CHLORPROMAZINE HYDROCHLORIDE

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	
DR REDDYS	25MG/ML	A080365 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	
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SYRUP; ORAL

CHLORPROMAZINE HYDROCHLORIDE

ALPHARMA US PHARMS	10MG/5ML	A086712 001	
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SONAZINE

FOSUN PHARMA	10MG/5ML	A083040 001	
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THORAZINE

+ GLAXOSMITHKLINE	10MG/5ML **	N009149 022	
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TABLET; ORAL

CHLORPROMAZINE HYDROCHLORIDE

ABBOTT	10MG	A084414 001	
	25MG	A084415 001	
	50MG	A084411 001	
	100MG	A084412 001	
	200MG	A084413 001	
CYCLE PHARMS LTD	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	
	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	
SANDOZ	10MG **	A080439 001	
	25MG **	A080439 002	
	50MG **	A080439 003	
	100MG **	A080439 004	

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

6-98(of 430)

** See List Footnote

CHLORPROMAZINE HYDROCHLORIDE

TABLET; ORAL

CHLORPROMAZINE HYDROCHLORIDE

	200MG **	A080439 005	
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 INC	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
AUROLIFE PHARMA LLC	100MG	A088725 001	Aug 31, 1984
	250MG	A088726 001	Aug 31, 1984
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
MYLAN	100MG	A088549 002	Jun 01, 1984
	250MG	A088549 001	Jun 01, 1984
PAR PHARM	100MG	A088175 001	Feb 27, 1984
	250MG	A088176 001	Feb 27, 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	

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

CHLORPROPAMIDETABLET; ORAL
GLUCAMIDE

ANI PHARMS INC 250MG A088641 001 Oct 11, 1984

CHLORPROTHIXENECONCENTRATE; 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 HYDROCHLORIDEOINTMENT; OPHTHALMIC
AUREOMYCIN

LEDERLE 1% N050404 001

CHLORTHALIDONE

TABLET; ORAL

CHLORTHALIDONE

ABBOTT 25MG A087364 001

50MG A087384 001

ACP NIMBLE 50MG A088651 001 May 30, 1985

ANI PHARMS INC 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

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

50MG A087118 001

50MG A087381 001

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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

CHLORTHALIDONE

TABLET; ORAL

THALITONE

+	CASPER PHARMA LLC	15MG **	N019574 001	Dec 20, 1988
		25MG	N019574 002	Feb 12, 1992
	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

	MYLAN	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	
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REGROTON

	SANOFI AVENTIS US	50MG; 0.25MG	N015103 001	
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CHLORZOXAZONE

TABLET; ORAL

CHLORZOXAZONE

	ACTAVIS ELIZABETH	250MG	A088928 001	May 08, 1987
		500MG	A040113 001	Sep 29, 1995
	AUROLIFE PHARMA LLC	250MG	A089852 001	May 04, 1988
		500MG	A089853 001	May 04, 1988
	BARR	500MG	A089895 001	May 04, 1988
	OHM LABS	250MG	A081298 001	Dec 29, 1993
		500MG	A081299 001	Dec 29, 1993
	PAR PHARM	250MG	A087981 001	Sep 20, 1983
	PIONEER PHARMS	250MG	A089592 001	Jan 06, 1989
		500MG	A089948 001	Jan 06, 1989
	SUN PHARM INDUSTRIES	500MG	A089970 001	Sep 27, 1990
	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	
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PARAFON FORTE DSC

+	JANSSEN R AND D	500MG **	N011529 002	Jun 15, 1987
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STRIFON FORTE DSC

	FERNDAL LABS	500MG	A081008 001	Dec 23, 1988
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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

	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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

CHOLESTYRAMINE

POWDER; ORAL

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

LOCHOLEST

ALLIED

EQ 4GM RESIN/PACKET

A074561 001 Aug 15, 1996

EQ 4GM RESIN/SCOOPFUL

A074561 002 Aug 15, 1996

LOCHOLEST LIGHT

ALLIED

EQ 4GM RESIN/PACKET

A074562 001 Aug 15, 1996

EQ 4GM RESIN/SCOOPFUL

A074562 002 Aug 15, 1996

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 FENOFIBRATE

CAPSULE, DELAYED RELEASE; ORAL

FENOFIBRIC ACID

MYLAN PHARMS INC

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

CHORIOGONADOTROPIN ALFA

INJECTABLE; INJECTION

OVIDREL

EMD SERONO

0.25MG/VIAL

N021149 001 Sep 20, 2000

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

CHYMOPAPAIN

INJECTABLE; INJECTION

CHYMODIACTIN

CHART MEDCL

4,000 UNITS/VIAL

N018663 002 Aug 21, 1984

+

10,000 UNITS/VIAL **

N018663 001 Nov 10, 1982

DISCASE

ABBOTT

12,500 UNITS/VIAL

N018625 001 Jan 18, 1984

CHYMOTRYPSIN

FOR SOLUTION; OPHTHALMIC

ALPHA CHYMAR

SOLA BARNES HIND

750 UNITS/VIAL

N011837 001

CATARASE

CIBA

300 UNITS/VIAL

N016938 001

NOVARTIS

150 UNITS/VIAL

N018121 001

ZOLYSE

ALCON

750 UNITS/VIAL

N011903 001

CICLOPIROX

SOLUTION; TOPICAL

CICLOPIROX

MYLAN PHARMS INC

8%

A078567 001 Sep 18, 2007

TEVA PHARMS

8%

A078079 001 Sep 18, 2007

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

CIDOFOVIR

INJECTABLE; INJECTION

VISTIDE

+ GILEAD SCIENCES INC EQ 75MG BASE/ML ** N020638 001 Jun 26, 1996

CILASTATIN SODIUM; IMPENEM

INJECTABLE; INJECTION

PRIMAXIN

MERC 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

MERC EQ 500MG BASE/VIAL; 500MG/VIAL N050630 001 Dec 14, 1990

EQ 750MG BASE/VIAL; 750MG/VIAL N050630 002 Dec 14, 1990

POWDER; INTRAVENOUS

IMPENEM 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

+ MERC EQ 250MG BASE/VIAL; 250MG/VIAL N050587 001 Nov 26, 1985

CILOSTAZOL

TABLET; ORAL

CILOSTAZOL

ACTAVIS ELIZABETH 100MG A077028 002 Nov 26, 2004

EPIC PHARMA LLC 50MG A077150 001 Mar 11, 2005

100MG A077022 001 Nov 23, 2004

IVAX SUB TEVA PHARMS 100MG A077020 002 Mar 01, 2005

MYLAN 50MG A077323 002 Apr 20, 2006

100MG A077323 001 Apr 20, 2006

MYLAN PHARMS INC 50MG A077019 001 Nov 23, 2004

100MG A077019 002 Nov 23, 2004

PLIVA HRVATSKA DOO 50MG A077898 001 Oct 29, 2007

100MG A077898 002 Oct 29, 2007

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 MOLECULES 200MG A074329 002 May 17, 1994

300MG A074329 003 May 17, 1994

400MG A074329 004 May 17, 1994

800MG A074329 001 May 17, 1994

CONTRACT PHARMACAL 200MG A074961 001 Jun 19, 1998

200MG A074963 001 Jun 19, 1998

CYCLE PHARMS LTD 300MG A074361 001 Dec 23, 1994

400MG A074361 002 Dec 23, 1994

800MG A074371 001 Dec 23, 1994

DAVA PHARMS INC 300MG A074340 001 Jun 23, 1995

400MG A074340 002 Jun 23, 1995

800MG A074339 001 Jun 23, 1995

IVAX SUB TEVA PHARMS 200MG A074401 001 May 30, 1995

200MG A074424 001 Jul 28, 1995

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

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

SANDOZ INC 100MG A075122 001 Jun 19, 1998

200MG A074250 001 Jun 29, 1995

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

CIMETIDINETABLET; ORAL
CIMETIDINE

	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
YAOPHARMA CO LTD	200MG		A074100	001	Jan 31, 1995
	300MG		A074100	002	Jan 31, 1995
	400MG		A074100	003	Jan 31, 1995
	800MG		A074100	004	Jan 31, 1995
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

CIMETIDINE HYDROCHLORIDE

INJECTABLE; INJECTION

CIMETIDINE HYDROCHLORIDE

HOSPIRA	EQ 300MG BASE/2ML		A074296	001	Mar 28, 1997
	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
VINTAGE PHARMS LLC	EQ 300MG BASE/2ML		A074005	001	Aug 31, 1994
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 INC	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 PHARMS LTD	EQ 300MG BASE/5ML		A074541	001	Aug 05, 1997
G AND W LABS INC	EQ 300MG BASE/5ML		A074176	001	Jun 01, 1994
LANNETT CO INC	EQ 300MG BASE/5ML		A074251	001	Dec 22, 1994
PHARM ASSOC	EQ 300MG BASE/5ML		A075560	001	Mar 15, 2000
TAGAMET					
GLAXOSMITHKLINE	EQ 300MG BASE/5ML **		N017924	001	

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

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

CINOXACIN

CAPSULE; ORAL

CINOAC

LILLY	250MG	N018067 001	
	500MG	N018067 002	

CINOXACIN

TEVA	250MG	A073005 001	Feb 28, 1992
	500MG	A073006 001	Feb 28, 1992

CIPROFLOXACIN

FOR SUSPENSION; ORAL

CIPROFLOXACIN

LUPIN LTD	250MG/5ML	A200563 001	Mar 05, 2014
	500MG/5ML	A200563 002	Mar 05, 2014

INJECTABLE; INJECTION

CIPRO

+ BAYER HLTHCARE	400MG/40ML (10MG/ML) **	N019847 001	Dec 26, 1990
+ BAYER HLTHCARE	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
+ BAYER HLTHCARE	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
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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

CIPROFLOXACIN IN DEXTROSE 5%

HIKMA FARMACEUTICA	200MG/100ML	A076757 001	Apr 21, 2008
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CIPROFLOXACIN IN DEXTROSE 5% IN PLASTIC CONTAINER

BAXTER HLTHCARE	200MG/100ML	A077888 001	Mar 18, 2008
	400MG/200ML	A077888 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

CIPROFLOXACIN HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

CIPROFLOXACIN HYDROCHLORIDE

AMRING PHARMS	EQ 0.3% BASE	A078598 001	Jan 16, 2008
RUBICON	EQ 0.3% BASE	A075928 001	Jun 09, 2004

TABLET; ORAL

CIPRO

+ BAYER HLTHCARE	EQ 100MG BASE **	N019537 001	Apr 08, 1996
+ BAYER HLTHCARE	EQ 750MG BASE **	N019537 004	Oct 22, 1987

CIPROFLOXACIN HYDROCHLORIDE

ANI PHARMS INC	EQ 100MG BASE	A075939 001	Mar 03, 2005
	EQ 250MG BASE	A075939 002	Jun 09, 2004
	EQ 500MG BASE	A075939 003	Jun 09, 2004
	EQ 750MG BASE	A075939 004	Jun 09, 2004
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
FOSUN PHARMA	EQ 250MG BASE	A076593 002	Jun 09, 2004
	EQ 500MG BASE	A076593 003	Jun 09, 2004
	EQ 750MG BASE	A076593 004	Jun 09, 2004
MYLAN	EQ 100MG BASE	A075817 001	Jun 25, 2007
	EQ 250MG BASE	A075685 002	Jun 09, 2004

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

CIPROFLOXACIN HYDROCHLORIDE

TABLET;ORAL

CIPROFLOXACIN HYDROCHLORIDE

	EQ 250MG BASE	A075817 002	Jun 09, 2004
	EQ 500MG BASE	A075685 003	Jun 09, 2004
	EQ 750MG BASE	A075685 001	Jun 09, 2004
	EQ 750MG BASE	A075817 004	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 100MG BASE	A076426 001	Jun 15, 2005
	EQ 250MG BASE	A076426 002	Jun 15, 2005
	EQ 500MG BASE	A076426 003	Jun 15, 2005
	EQ 750MG BASE	A076426 004	Jun 15, 2005
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
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

TABLET, EXTENDED RELEASE;ORAL

PROQUIN XR

DEPOMED INC	EQ 500MG BASE	N021744 001	May 19, 2005
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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 INC	212.6MG;EQ 287.5MG BASE	A077417 001	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
FOSUN PHARMA	212.6MG;EQ 287.5MG BASE	A078712 001	Dec 11, 2007
MYLAN PHARMS INC	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
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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
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CISATRACURIUM BESYLATE

INJECTABLE;INJECTION

CISATRACURIUM BESYLATE

ACCORD HLTHCARE	EQ 2MG BASE/ML	A205873 001	Jun 16, 2017
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CISATRACURIUM BESYLATE PRESERVATIVE FREE

ACCORD HLTHCARE	EQ 2MG BASE/ML	A205872 001	Jun 16, 2017
	EQ 10MG BASE/ML	A205872 002	Jun 16, 2017

CISPLATIN

INJECTABLE;INJECTION

CISPLATIN

BEDFORD	10MG/VIAL	A074713 001	Nov 14, 2000
	50MG/VIAL	A074713 002	Nov 14, 2000
MYLAN LABS LTD	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
+		50MG/VIAL	N018057 002

PLATINOL-AQ

+	HQ SPCLT PHARMA	0.5MG/ML	N018057 003	Jul 18, 1984
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Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** 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
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CLADRIBINE

INJECTABLE;INJECTION

LEUSTATIN

+	JANSSEN PHARMS	1MG/ML **	N020229 001	Feb 26, 1993
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CLARITHROMYCIN

FOR SUSPENSION;ORAL

BIAXIN

+	ABBVIE	125MG/5ML	N050698 001	Dec 23, 1993
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		187MG/5ML	N050698 003	Sep 30, 1998
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+		250MG/5ML	N050698 002	Dec 23, 1993
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CLARITHROMYCIN

SUN PHARM INDS LTD	125MG/5ML	A065382 001	Aug 30, 2007
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	250MG/5ML	A065382 002	Aug 30, 2007
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TABLET;ORAL

BIAXIN

+	ABBVIE	250MG **	N050662 001	Oct 31, 1991
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+		500MG **	N050662 002	Oct 31, 1991
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CLARITHROMYCIN

AJANTA PHARMA LTD	250MG	A206714 001	Apr 25, 2019
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	500MG	A206714 002	Apr 25, 2019
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IVAX SUB TEVA PHARMS	250MG	A065137 001	May 31, 2005
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	500MG	A065137 002	May 31, 2005
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MYLAN	250MG	A065195 001	Mar 11, 2005
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	500MG	A065195 002	Mar 11, 2005
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SUN PHARM INDS LTD	250MG	A065174 001	Sep 24, 2004
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	500MG	A065174 002	Sep 24, 2004
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TABLET, EXTENDED RELEASE;ORAL

BIAXIN XL

+	ABBVIE	500MG **	N050775 001	Mar 03, 2000
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CLARITHROMYCIN

ANI PHARMS INC	500MG	A065250 001	Aug 25, 2005
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LUPIN LTD	500MG	A202532 001	Sep 15, 2015
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RANBAXY	1GM	A065210 001	Jan 26, 2005
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CLAVULANATE POTASSIUM; TICARCILLIN DISODIUM

INJECTABLE;INJECTION

TIMENTIN

GLAXOSMITHKLINE	EQ 100MG BASE/VIAL;EQ 3GM BASE/VIAL	A062691 001	Dec 19, 1986
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	EQ 100MG BASE/VIAL;EQ 3GM BASE/VIAL	N050590 001	Apr 01, 1985
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	EQ 200MG BASE/VIAL;EQ 3GM BASE/VIAL	N050590 002	Apr 01, 1985
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	EQ 1GM BASE/VIAL;EQ 30GM BASE/VIAL	N050590 003	Aug 18, 1987
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TIMENTIN IN PLASTIC CONTAINER

GLAXOSMITHKLINE	EQ 100MG BASE/100ML;EQ 3GM BASE/100ML	N050658 001	Dec 15, 1989
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CLEMASTINE FUMARATE

SYRUP;ORAL

CLEMASTINE FUMARATE

ACTAVIS MID ATLANTIC	EQ 0.5MG BASE/5ML	A074075 001	Oct 31, 1993
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APOTEX INC	EQ 0.5MG BASE/5ML	A075703 001	Nov 27, 2000
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LANNETT CO INC	EQ 0.5MG BASE/5ML	A074884 001	Dec 17, 1997
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TEVA PHARMS	EQ 0.5MG BASE/5ML	A073095 001	Apr 21, 1992
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WOCKHARDT BIO AG	EQ 0.5MG BASE/5ML	A074863 001	Mar 13, 1998
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TAVIST

+	NOVARTIS	EQ 0.5MG BASE/5ML **	N018675 001	Jun 28, 1985
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TABLET;ORAL

CLEMASTINE FUMARATE

ANI PHARMS INC	1.34MG	A073282 001	Jan 31, 1992
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	1.34MG	A073282 002	Dec 03, 1992
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PLD ACQUISITIONS LLC	1.34MG	A073458 001	Oct 31, 1993
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SANDOZ	2.68MG	A073459 001	Oct 31, 1993
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TAVIST

+	NOVARTIS	2.68MG	N017661 001
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TAVIST-1

+	GLAXOSMITHKLINE CONS	1.34MG	N020925 001	Aug 21, 1992
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NOVARTIS	1.34MG	N017661 002	
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Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

CLEMASTINE FUMARATETABLET; ORAL
TAVIST-1

1.34MG

N017661 003 Aug 21, 1992

CLEVIDIPINEEMULSION; INTRAVENOUS
CLEVIPREX

+ CHIESI USA INC

125MG/250ML (0.5MG/ML)

N022156 003 Nov 08, 2013

CLIDINIUM BROMIDECAPSULE; 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

MYLAN PHARMS INC

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

CLINDAMYCIN PALMITATE HYDROCHLORIDE

FOR SOLUTION; ORAL

CLEOCIN

PHARMACIA AND UPJOHN

EQ 75MG BASE/5ML **

A061827 001

CLINDAMYCIN PALMITATE HYDROCHLORIDE

MYLAN

EQ 75MG BASE/5ML

A203063 001 May 25, 2016

CLINDAMYCIN PHOSPHATE

CREAM; VAGINAL

CLEOCIN

PHARMACIA AND UPJOHN

EQ 2% BASE

N050680 001 Aug 11, 1992

INJECTABLE; INJECTION

CLEOCIN PHOSPHATE

PHARMACIA AND UPJOHN

EQ 150MG BASE/ML

A061839 001

CLINDAMYCIN PHOSPHATE

ABRAXIS PHARM

EQ 150MG BASE/ML

A062747 001 Jun 03, 1988

BEDFORD

EQ 150MG BASE/ML

A063163 001 Jun 30, 1994

BRISTOL MYERS SQUIBB

EQ 150MG BASE/ML

A062908 001 Feb 01, 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

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

WEST-WARD PHARMS INT

EQ 150MG BASE/ML

A062806 001 Oct 15, 1987

EQ 150MG BASE/ML

A062953 001 Apr 21, 1988

EQ 150MG BASE/ML

A063068 001 Aug 28, 1989

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

PHARMACIA AND UPJOHN

EQ 1% BASE

A062363 001 Feb 08, 1982

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

CLINDAMYCIN PHOSPHATE

SOLUTION; TOPICAL

CLINDAMYCIN PHOSPHATE

BOCA PHARMA LLC	EQ 1% BASE	A062944 001	Jan 11, 1989
NOVAST LABS	EQ 1% BASE	A064108 001	Sep 27, 1996
VINTAGE PHARMS	EQ 1% BASE	A062930 001	Jun 28, 1989
WOCKHARDT BIO AG	EQ 1% BASE	A063304 001	Jul 15, 1997

CLIOQUINOL; NYSTATIN

OINTMENT; TOPICAL

NYSTAFORM

BAYER PHARMS	10MG/GM;100,000 UNITS/GM	N050235 001	
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CLOBAZAM

SUSPENSION; ORAL

CLOBAZAM

TARO	2.5MG/ML	A210978 001	Apr 15, 2019
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TABLET; ORAL

CLOBAZAM

ACCORD HLTHCARE	10MG	A212398 001	May 23, 2019
	20MG	A212398 002	May 23, 2019
TARO	10MG	A209440 001	Oct 22, 2018
	20MG	A209440 002	Oct 22, 2018

ONFI

LUNDBECK PHARMS LLC	5MG **	N202067 001	Oct 21, 2011
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CLOBETASOL PROPIONATE

CREAM; TOPICAL

CLOBETASOL PROPIONATE

TEVA PHARMS USA	0.05%	A074087 001	Feb 16, 1994
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TEMOVATE

+ FOUGERA PHARMS	0.05% **	N019322 001	Dec 27, 1985
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TEMOVATE E

+ FOUGERA PHARMS	0.05% **	N020340 001	Jun 17, 1994
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GEL; TOPICAL

TEMOVATE

+ FOUGERA PHARMS	0.05% **	N020337 001	Apr 29, 1994
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LOTION; TOPICAL

CLOBETASOL PROPIONATE

HI TECH	0.05%	A211348 001	Oct 26, 2018
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OINTMENT; TOPICAL

CLOBETASOL PROPIONATE

ACTAVIS MID ATLANTIC	0.05%	A074128 001	Aug 03, 1994
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AMNEAL PHARMS LLC	0.05%	A210551 001	Aug 21, 2018
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TEMOVATE

+ FOUGERA PHARMS	0.05% **	N019323 001	Dec 27, 1985
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SOLUTION; TOPICAL

TEMOVATE

+ FOUGERA PHARMS	0.05% **	N019966 001	Feb 22, 1990
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SPRAY; TOPICAL

CLOBETASOL PROPIONATE

ALEOR DERMACEUTICALS	0.05%	A211191 001	Oct 02, 2019
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APOTEX	0.05%	A210446 001	Apr 17, 2018
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CLOFARABINE

SOLUTION; INTRAVENOUS

CLOFARABINE

HOSPIRA INC	20MG/20ML (1MG/ML)	A210283 001	Dec 27, 2018
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CLOFAZIMINE

CAPSULE; ORAL

LAMPRENE

+ NOVARTIS	50MG	N019500 002	Dec 15, 1986
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	100MG	N019500 001	Dec 15, 1986
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CLOFIBRATE

CAPSULE; ORAL

ATROMID-S

WYETH AYERST	500MG	N016099 002	
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CLOFIBRATE

BANNER PHARMACAPS	500MG	A073396 001	Mar 20, 1992
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SANDOZ	500MG	A072191 001	May 02, 1988
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Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

6-110(of 430)

** See List Footnote

CLOFIBRATE

CAPSULE; ORAL

CLOFIBRATE

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

MILEX 50MG

A072196 001 Dec 20, 1988

SEROPHENE

EMD SERONO 50MG

N018361 001 Mar 22, 1982

CLOMIPRAMINE HYDROCHLORIDE

CAPSULE; ORAL

CLOMIPRAMINE HYDROCHLORIDE

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

CLONAZEPAM

TABLET; ORAL

CLONAZEPAM

MYLAN PHARMS INC	0.5MG	A074940	001	Oct 30, 1997
	1MG	A074940	002	Oct 30, 1997
	2MG	A074940	003	Oct 30, 1997
RUBICON	0.5MG	A075468	001	Oct 06, 2000
	1MG	A075468	002	Oct 06, 2000
	2MG	A075468	003	Oct 06, 2000
SANDOZ	0.5MG	A074925	001	Sep 30, 1997
	1MG	A074925	002	Sep 30, 1997
	2MG	A074925	003	Sep 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
KLONOPIN				
ROCHE	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
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TABLET, EXTENDED RELEASE; ORAL

CLONIDINE

TRIS PHARMA INC	EQ 0.17MG BASE	N022500	001	Dec 03, 2009
	EQ 0.26MG BASE	N022500	002	Dec 03, 2009

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

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

TABLET; ORAL

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
AUROLIFE PHARMA LLC	0.1MG	A070886 002	Aug 31, 1988
	0.2MG	A070886 001	Aug 31, 1988
	0.3MG	A070886 003	Aug 31, 1988
CHARTWELL MOLECULES	0.1MG	A071785 002	Apr 05, 1988
	0.2MG	A071785 003	Apr 05, 1988
	0.3MG	A071785 001	Apr 05, 1988
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
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
	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	0.2MG	A202792 002	May 15, 2015
	0.2MG	A203320 002	May 15, 2015
ANCHEN PHARMS	0.1MG	A202983 001	Apr 02, 2014
	0.2MG	A202983 002	Apr 02, 2014
	0.2MG	A202984 002	Sep 30, 2013

JENLOGA

+ CONCORDIA PHARMS INC	0.1MG **	N022331 001	Sep 30, 2009
+ CONCORDIA PHARMS INC	0.2MG **	N022331 002	May 25, 2010

KAPVAY

+ CONCORDIA PHARMS INC	0.2MG **	N022331 004	Sep 28, 2010
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CLOPIDOGREL BISULFATE

TABLET; ORAL

CLOPIDOGREL BISULFATE

ACTAVIS TOTOWA	EQ 75MG BASE	A090307 001	May 28, 2013
ANI PHARMS INC	EQ 300MG BASE	A090625 001	May 17, 2012
	EQ 300MG BASE	A091216 001	May 17, 2012
CADILA	EQ 75MG BASE	A201686 001	Oct 10, 2012
MYLAN PHARMS INC	EQ 75MG BASE	A077665 001	May 17, 2012
	EQ 300MG BASE	A077665 002	May 17, 2012
SUN PHARM INDUSTRIES	EQ 75MG BASE	A078133 001	Jun 10, 2013

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

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
AUROLIFE PHARMA LLC	3.75MG	A072112	002	Aug 26, 1988
	7.5MG	A072112	003	Aug 26, 1988
	15MG	A072112	001	Aug 26, 1988
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
MYLAN	3.75MG	A071509	001	Oct 19, 1987
	7.5MG	A071510	001	Oct 19, 1987
	15MG	A071511	001	Oct 19, 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
	7.5MG	A071550	001	Sep 12, 1988
	15MG	A071522	001	Sep 12, 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				
RECORDATI RARE	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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

CLORAZEPATE DIPOTASSIUM

TABLET;ORAL

TRANXENE

RECORDATI RARE 3.75MG ** N017105 006

+ 15MG ** N017105 008

TRANXENE SD

RECORDATI RARE 11.25MG ** N017105 005

22.5MG ** N017105 004

CLOTRIMAZOLE

CREAM;TOPICAL

LOTRIMIN

SCHERING PLOUGH 1% ** N017619 001

MYCELEX

BAYER HEALTHCARE LLC 1% N018183 001

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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

CLOXACILLIN SODIUMFOR SOLUTION;ORAL
TEGOPEN

EQ 125MG BASE/5ML N050192 001

CLOZAPINETABLET;ORAL
CLOZAPINE

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
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

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

COBALT CHLORIDE CO-57; CYANOCOBALAMIN; CYANOCOBALAMIN CO-57; INTRINSIC FACTOR

N/A;N/A

RUBRATOPE-57 KIT

BRACCO N/A;N/A;N/A;N/A N016089 001

COBALT CHLORIDE CO-60; CYANOCOBALAMIN; CYANOCOBALAMIN CO-60; INTRINSIC FACTOR

N/A;N/A

RUBRATOPE-60 KIT

BRACCO N/A;N/A;N/A;N/A N016090 001

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

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

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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

COLCHICINE

CAPSULE; ORAL			
COLCHICINE			
PAR PHARM INC	0.6MG	A208678 001	Nov 29, 2018
TABLET; ORAL			
COLCHICINE			
ZYDUS PHARMS	0.6MG	A211519 001	Feb 19, 2019

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 INC	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

CAPSULE; ORAL			
WELCHOL			
DAIICHI SANKYO	375MG	N021141 001	May 26, 2000
FOR SUSPENSION; ORAL			
WELCHOL			
+ DAIICHI SANKYO	1.875GM/PACKET	N022362 001	Oct 02, 2009

COLISTIN SULFATE

SUSPENSION; ORAL			
COLY-MYCIN S			
PARKE DAVIS	EQ 25MG BASE/5ML	N050355 001	

CONIVAPTAN HYDROCHLORIDE

INJECTABLE; INTRAVENOUS			
VAPRISOL			
CUMBERLAND PHARMS	20MG/4ML (5MG/ML)	N021697 001	Dec 29, 2005

COPPER

INTRAUTERINE DEVICE; INTRAUTERINE			
CU-7			
GD SEARLE LLC	89MG	N017408 001	
TATUM-T			
GD SEARLE LLC	120MG	N018205 001	

CORTICOTROPIN

INJECTABLE; INJECTION			
ACTH			
PARKEDALE	25 UNITS/VIAL	N008317 002	
	40 UNITS/VIAL	N008317 004	
ACTHAR			
SANOFI AVENTIS US	25 UNITS/VIAL	N007504 002	
	40 UNITS/VIAL	N007504 003	
CORTICOTROPIN			
ORGANICS LAGRANGE	40 UNITS/ML	N010831 001	
	80 UNITS/ML	N010831 002	
WATSON LABS	40 UNITS/VIAL	A088772 001	Nov 21, 1984
H.P. ACTHAR GEL			
MALLINCKRODT ARD	40 UNITS/ML	N008372 006	
PURIFIED CORTROPHIN GEL			
ANI PHARMS INC	40 UNITS/ML	N008975 001	
	80 UNITS/ML	N008975 002	

CORTICOTROPIN-ZINC HYDROXIDE

INJECTABLE; INJECTION			
CORTROPHIN-ZINC			
ANI PHARMS INC	40 UNITS/ML	N009854 001	

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

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
ELKINS SINN	25MG	A080836 001
EVERYLIFE	25MG	A084246 001
HEATHER	25MG	A085736 001
IMPAX LABS	25MG	N009458 001
INWOOD LABS	25MG	A080731 001
IVAX SUB TEVA PHARMS	25MG	A080630 001
	25MG	A083536 001
LANNETT	25MG	A080694 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
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COSYNTROPIN

SOLUTION; INTRAVENOUS

COSYNTROPIN

SANDOZ INC	0.25MG/ML (0.25MG/ML)	N022028 001	Feb 21, 2008
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CROMOLYN SODIUM

AEROSOL, METERED; INHALATION

INTAL

KING PHARMS LLC	0.8MG/INH	N018887 001	Dec 05, 1985
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CAPSULE; INHALATION

INTAL

+ SANOFI AVENTIS US	20MG **	N016990 001
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CAPSULE; ORAL

GASTROCROM

UCB INC	100MG	N019188 001	Dec 22, 1989
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CONCENTRATE; ORAL

CROMOLYN SODIUM

GENERA PHARMS	100MG/5ML	A090954 001	Dec 18, 2009
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SOLUTION; INHALATION

CROMOLYN SODIUM

ACTAVIS MID ATLANTIC	10MG/ML	A075067 001	Jul 19, 1999
BAUSCH AND LOMB	10MG/ML	A075585 001	Dec 21, 2000
FERA PHARMS LLC	10MG/ML	A075437 001	Apr 21, 2000
HIKMA	10MG/ML	A075333 001	Apr 30, 2002
MYLAN SPECIALITY LP	10MG/ML	A074209 001	Apr 26, 1994
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
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SOLUTION/DROPS; OPHTHALMIC

CROLOM

BAUSCH AND LOMB	4%	A074443 001	Jan 30, 1995
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CROMOLYN SODIUM

APOTEX INC	4%	A075615 001	Jan 26, 2001
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CROMOPTIC

KING PHARMS	4%	A075088 001	Apr 27, 1999
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Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

6-117(of 430)

** See List Footnote

CROMOLYN SODIUMSOLUTION/DROPS;OPHTHALMIC
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

NASALCROM

+ BLACKSMITH BRANDS 5.2MG/SPRAY **

N020463 001 Jan 03, 1997

CROTAMITON

CREAM;TOPICAL

EURAX

+ SUN PHARM INDS INC 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

PAR PHARM 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

AKORN 1MG/ML

A087969 001 Nov 10, 1983

DELL LABS 0.03MG/ML

A080689 001

0.1MG/ML

A080689 002

1MG/ML

A080689 003

DR REDDYS 0.1MG/ML

A080573 002

1MG/ML

A080573 001

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

DODEX

ACCORD HLTHCARE 1MG/ML

A083022 001

REDISOL

MERCK 1MG/ML

N006668 010

RUBIVITE

BEL MAR 0.03MG/ML

N010791 004

0.05MG/ML

N010791 001

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

6-118(of 430)

** See List Footnote

CYANOCOBALAMIN

INJECTABLE; INJECTION

RUBIVITE

0.1MG/ML	N010791 002
0.12MG/ML	N010791 005
1MG/ML	N010791 003

RUBRAMIN PC

BRISTOL MYERS SQUIBB	0.1MG/ML	N006799 002
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+	1MG/ML **	N006799 004
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+	1MG/ML **	N006799 010	Apr 28, 1988
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RUVITE

SAVAGE LABS	1MG/ML	A080570 002
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VI-TWEL

BAYER HLTHCARE	1MG/ML	N007012 002
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SPRAY, METERED; NASAL

CALOMIST

PAR PHARM	25MCG/SPRAY	N022102 001	Jul 27, 2007
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TABLET; ORAL

CYANOCOBALAMIN

WEST WARD	1MG	A084264 001
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CYANOCOBALAMIN CO-57

CAPSULE; ORAL

RUBRATOPE-57

BRACCO	0.5-1uCi	N016089 002
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CYANOCOBALAMIN CO-60

CAPSULE; ORAL

RUBRATOPE-60

BRACCO	0.5-1uCi	N016090 002
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CYANOCOBALAMIN; CYANOCOBALAMIN CO-57; CYANOCOBALAMIN CO-58

N/A; N/A

DICOPAC KIT

GE HEALTHCARE	N/A; N/A; N/A	N017406 001
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CYANOCOBALAMIN; CYANOCOBALAMIN CO-57; INTRINSIC FACTOR

N/A; N/A

CYANOCOBALAMIN CO 57 SCHILLING TEST KIT

MALLINCKRODT	0.1MG; 0.5uCi; 60MG	N016635 001
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CYANOCOBALAMIN; TANNIC ACID; ZINC ACETATE

INJECTABLE; INJECTION

DEPINAR

ARMOUR PHARM	0.5MG/ML; 2.3MG/ML; 1MG/ML	N011208 001
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CYCLACILLIN

FOR SUSPENSION; ORAL

CYCLAPEN-W

WYETH AYERST	125MG/5ML	N050508 001
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	250MG/5ML	N050508 002
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	500MG/5ML	N050508 003
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TABLET; ORAL

CYCLACILLIN

TEVA	250MG	A062895 001	Aug 04, 1988
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	500MG	A062895 002	Aug 04, 1988
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CYCLAPEN-W

WYETH AYERST	250MG	N050509 001
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	500MG	N050509 002
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CYCLIZINE LACTATE

INJECTABLE; INJECTION

MAREZINE

GLAXOSMITHKLINE	50MG/ML	N009495 001
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CYCLOBENZAPRINE HYDROCHLORIDE

TABLET; ORAL

CYCLOBENZAPRINE HYDROCHLORIDE

SANDOZ	10MG	A073683 001	Feb 26, 1993
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UPSHER SMITH LABS	5MG	A072854 002	Feb 03, 2006
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	10MG	A072854 001	Nov 19, 1991
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WATSON LABS	10MG	A073143 001	Nov 27, 1991
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	10MG	A074436 001	Nov 30, 1994
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Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

6-119(of 430)

** See List Footnote

CYCLOBENZAPRINE HYDROCHLORIDE

TABLET; ORAL

FLEXERIL

+ JANSSEN RES AND DEV	5MG **	N017821 001	
+	10MG **	N017821 002	

CYCLOPENTOLATE HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

AK-PENTOLATE

AKORN	1%	A085555 001	
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AKPENTOLATE

AKORN	2%	A040165 001	Jan 13, 1997
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CYCLOPENTOLATE HYDROCHLORIDE

ALCON PHARMS LTD	1%	A089162 001	Jan 24, 1991
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SOLA BARNES HIND	1%	A084150 001	
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	1%	A084863 001	
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PENTOLAIR

PHARMAFAIR	0.5%	A088643 001	Feb 09, 1987
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	1%	A088150 001	Feb 25, 1983
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CYCLOPHOSPHAMIDE

INJECTABLE; INJECTION

CYCLOPHOSPHAMIDE

BAXTER HLTHCARE	100MG/VIAL	A088371 001	Jul 03, 1986
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	200MG/VIAL	A088372 001	Jul 03, 1986
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	500MG/VIAL	A088373 001	Jul 03, 1986
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	1GM/VIAL	A088374 001	Sep 24, 1986
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CYTOXAN

+ BAXTER HLTHCARE	100MG/VIAL **	N012142 001	
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+	200MG/VIAL **	N012142 002	
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CYTOXAN (LYOPHILIZED)

+ BAXTER HLTHCARE	500MG/VIAL	N012142 003	
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+	500MG/VIAL **	N012142 008	Jan 04, 1984
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+	1GM/VIAL	N012142 004	Aug 30, 1982
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+	1GM/VIAL **	N012142 010	Sep 24, 1985
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+	2GM/VIAL	N012142 005	Aug 30, 1982
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+	2GM/VIAL **	N012142 009	Dec 10, 1985
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LYOPHILIZED CYTOXAN

+ BAXTER HLTHCARE	100MG/VIAL **	N012142 006	Dec 05, 1985
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+	200MG/VIAL **	N012142 007	Dec 10, 1985
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NEOSAR

BEDFORD	100MG/VIAL	A087442 001	Feb 16, 1982
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	200MG/VIAL	A087442 002	Feb 16, 1982
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	500MG/VIAL	A087442 003	Feb 16, 1982
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	1GM/VIAL	A087442 004	Jul 08, 1983
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	2GM/VIAL	A087442 005	Mar 30, 1989
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TEVA PARENTERAL	100MG/VIAL	A040015 001	Apr 29, 1993
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	200MG/VIAL	A040015 002	Apr 29, 1993
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	500MG/VIAL	A040015 003	Apr 29, 1993
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	1GM/VIAL	A040015 004	Apr 29, 1993
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	2GM/VIAL	A040015 005	Apr 29, 1993
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TABLET; ORAL

CYCLOPHOSPHAMIDE

ROXANE	25MG	A040032 001	Aug 17, 1999
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	50MG	A040032 002	Aug 17, 1999
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CYTOXAN

+ BAXTER HLTHCARE	25MG **	N012141 002	
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+	50MG **	N012141 001	
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CYCLOSPORINE

CAPSULE; ORAL

GENGRAF

ABBVIE	50MG	A065003 002	May 12, 2000
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NEORAL

+ NOVARTIS	50MG **	N050715 003	Jul 14, 1995
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SOLUTION; ORAL

CYCLOSPORINE

PHARM ASSOC	100MG/ML	A065167 001	Jan 05, 2005
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DISCONTINUED DRUG PRODUCT LIST

6-120(of 430)

** See List Footnote

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

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 A088212 001 May 26, 1983

DURAMED PHARMS BARR 4MG A088232 001 Oct 25, 1983

FOSUN PHARMA 4MG A086808 001

HALSEY 4MG A089057 001 Jul 03, 1986

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

SUPERPHARM 4MG A087405 001

VITARINE 4MG A087284 001

WATSON LABS 4MG A085245 001

4MG A086165 001

4MG A086580 001

PERIACTIN

+ MERCK 4MG ** N012649 001

CYSTEINE HYDROCHLORIDE

INJECTABLE; INJECTION

CYSTEINE HYDROCHLORIDE

+ HOSPIRA 7.25% ** N019523 001 Oct 22, 1986

CYTARABINE

INJECTABLE; INJECTION

CYTARABINE

MYLAN LABS LTD 20MG/ML A200916 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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

6-121(of 430)

** See List Footnote

DACARBAZINE

INJECTABLE; INJECTION

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

DACTINOMYCIN

	AM REGENT	0.5MG/VIAL	A202562 001	Aug 23, 2013
	WEST-WARD PHARMS INT	0.5MG/VIAL	A090304 001	Mar 16, 2010

DALFOPRISTIN; QUINUPRISTIN

INJECTABLE; INTRAVENOUS

SYNERCID

	KING PHARMS	420MG/VIAL; 180MG/VIAL	N050748 002	Aug 24, 2000
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DALTEPARIN SODIUM

INJECTABLE; INJECTION

FRAGMIN

	PFIZER INC	7,500 IU/0.75ML	N020287 008	Apr 04, 2002
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INJECTABLE; SUBCUTANEOUS

FRAGMIN

	PFIZER INC	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
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DANAZOL

CAPSULE; ORAL

DANAZOL

	AM THERAP	200MG	A071569 001	Dec 30, 1987
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DANOCRINE

	SANOFI AVENTIS US	50MG **	N017557 003
		100MG **	N017557 004
		200MG **	N017557 002

DANTROLENE SODIUM

INJECTABLE; INJECTION

DANTROLENE SODIUM

	MYLAN INSTITUTIONAL	20MG/VIAL	A205239 001	Feb 18, 2016
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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

DAPSONE

TABLET; ORAL

DAPSONE

	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
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DISCONTINUED DRUG PRODUCT LIST

6-122(of 430)

** See List Footnote

DARIFENACIN HYDROBROMIDE

TABLET, EXTENDED RELEASE;ORAL

DARIFENACIN HYDROBROMIDE

ANCHEN PHARMS

EQ 7.5MG BASE

A091190 001 Mar 13, 2015

EQ 15MG BASE

A091190 002 Mar 13, 2015

DARUNAVIR

TABLET;ORAL

PREZISTA

+ JANSSEN PRODS

300MG **

N021976 001 Jun 23, 2006

+

400MG **

N021976 003 Oct 21, 2008

DASABUVIR SODIUM; OMBITASVIR; PARITAPREVIR; RITONAVIR

TABLET, EXTENDED RELEASE;ORAL

VIEKIRA XR

+ ABBVIE INC

EQ 200MG BASE;8.33MG;50MG;33.33MG

N208624 001 Jul 22, 2016

DASATINIB

TABLET;ORAL

DASATINIB

APOTEX INC

20MG

A202103 001 Jun 10, 2016

50MG

A202103 002 Jun 10, 2016

70MG

A202103 003 Jun 10, 2016

100MG

A202103 004 Jun 10, 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

DEFERIPRONE

TABLET;ORAL

DEFERIPRONE

TARO PHARM INDS LTD

500MG

A208800 001 Feb 08, 2019

DEFEROXAMINE MESYLATE

INJECTABLE;INJECTION

DEFEROXAMINE MESYLATE

DR REDDYS

500MG/VIAL

A076806 001 Mar 31, 2006

2GM/VIAL

A076806 002 Mar 31, 2006

DELAVIRDINE MESYLATE

TABLET;ORAL

RESCRIPTOR

+ VIIV HLTHCARE

100MG

N020705 001 Apr 04, 1997

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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

6-123(of 430)

** See List Footnote

DEMECLOCYCLINE HYDROCHLORIDE

TABLET; ORAL

DECLOMYCIN

COREPHARMA	75MG	N050261 001
	150MG	N050261 002
	300MG	N050261 003

DEMECLOCYCLINE HYDROCHLORIDE

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
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ORETICYL 50

ABBVIE	0.125MG; 50MG	N012148 003
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ORETICYL FORTE

ABBVIE	0.25MG; 25MG	N012148 002
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DESERPIDINE; METHYCLOTHIAZIDE

TABLET; ORAL

ENDURONYL

ABBOTT	0.25MG; 5MG	N012775 001
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ENDURONYL FORTE

ABBOTT	0.5MG; 5MG	N012775 002
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METHYCLOTHIAZIDE 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 INC	25MG	A071803 002	Dec 08, 1987
	50MG	A071803 003	Dec 08, 1987
	75MG	A071803 004	Dec 08, 1987
	100MG	A071803 001	May 29, 1997
	150MG	A071803 005	May 29, 1997
USL PHARMA	25MG	A071864 001	Sep 09, 1987
	50MG	A071865 001	Sep 09, 1987
	75MG	A071866 001	Sep 09, 1987
	100MG	A071867 001	Sep 09, 1987

DESIRUDIN RECOMBINANT

INJECTABLE; SUBCUTANEOUS

IPRIVASK

+ BAUSCH	15MG/VIAL	N021271 001	Apr 04, 2003
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DESLANOSIDE

INJECTABLE; INJECTION

CEDILANID-D

NOVARTIS	0.2MG/ML	N009282 002
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DESLORATADINE

TABLET; ORAL

DESLORATADINE

MYLAN PHARMS INC	5MG	A078351 001	Feb 10, 2012
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DISCONTINUED DRUG PRODUCT LIST

6-124(of 430)

** See List Footnote

DESMOPRESSIN ACETATE

INJECTABLE; INJECTION

DDAVP

FERRING PHARMS INC 0.015MG/ML N018938 002 Apr 25, 1995

DESMOPRESSIN ACETATE

BEDFORD 0.004MG/ML A074575 001 Feb 18, 2000

HOSPIRA 0.004MG/ML A075220 001 Aug 28, 2000

DESMOPRESSIN ACETATE PRESERVATIVE FREE

BEDFORD 0.004MG/ML A074574 001 Feb 18, 2000

SOLUTION; NASAL

CONCENTRAID

FERRING 0.01% N019776 001 Dec 26, 1990

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

NOCTIVA

+ ALYVANT 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

TABLET; ORAL

DESMOPRESSIN ACETATE

FERRING 0.1MG N021795 001 May 08, 2008

0.2MG N021795 002 May 08, 2008

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

ACCORD HLTHCARE 0.15MG;0.03MG A207067 001 Sep 13, 2018

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

DESOXIMETASONE

CREAM; TOPICAL

DESOXIMETASONE

CADILA 0.25% A205620 001 Sep 28, 2018

TOPICORT

+ TARO PHARM INDS LTD 0.25% ** N017856 001

TOPICORT LP

+ TARO PHARM INDS LTD 0.05% ** N018309 001

GEL; TOPICAL

TOPICORT

+ TARO PHARM INDS LTD 0.05% ** N018586 001 Mar 29, 1982

OINTMENT; TOPICAL

DESOXIMETASONE

ALTANA 0.25% A073440 001 Apr 01, 1998

TOPICORT

+ TARO PHARM INDS LTD 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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

6-125(of 430)

** See List Footnote

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 PHARMA GLOBAL

EQ 50MG BASE

N205583 001 Jan 28, 2014

+

EQ 100MG BASE

N205583 002 Jan 28, 2014

TEVA PHARMS USA

EQ 50MG BASE

N205208 001 Oct 11, 2013

EQ 100MG BASE

N205208 002 Oct 11, 2013

DESVENLAFAXINE SUCCINATE

TABLET, EXTENDED RELEASE; ORAL

DESVENLAFAXINE SUCCINATE

MYLAN

EQ 50MG BASE

A204095 001 Jun 29, 2015

EQ 100MG BASE

A204095 002 Jun 29, 2015

YICHANG HUMANWELL

EQ 50MG BASE

A210014 001 Oct 01, 2018

EQ 100MG BASE

A210014 002 Oct 01, 2018

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

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

ANI PHARMS INC

0.75MG

A080399 001

HERITAGE PHARMA

1.5MG

A085456 001

IMPAX LABS

0.75MG

A085376 001

PAR PHARM

0.25MG

A088149 001 Apr 28, 1983

PHOENIX LABS NY

0.75MG

A083806 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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

6-126(of 430)

** See List Footnote

DEXAMETHASONE

TABLET; ORAL

DEXAMETHASONE

	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
WHITEWORTH 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

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

AKORN

EQ 4MG PHOSPHATE/ML

A084493 001

AUROBINDO PHARMA LTD

EQ 10MG PHOSPHATE/ML

A210967 001 Jun 07, 2019

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

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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

6-127(of 430)

** See List Footnote

DEXAMETHASONE SODIUM PHOSPHATE

INJECTABLE; INJECTION

HEXADROL

+

EQ 10MG PHOSPHATE/ML **

N014694 003

EQ 20MG PHOSPHATE/ML

N014694 004

OINTMENT; OPHTHALMIC

DECADRON

MERCCK

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

+ MERCCK

EQ 0.1% PHOSPHATE

N011984 001

DEXAMETHASONE SODIUM PHOSPHATE

SANDOZ INC

EQ 0.1% PHOSPHATE

A088771 001 Jan 16, 1985

SOLUTION/DROPS; OTIC

DEXAMETHASONE SODIUM PHOSPHATE

AKORN

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

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

DEXBROMPHENIRAMINE MALEATE

SYRUP; ORAL

DISOMER

SCHERING

2MG/5ML

N011814 002

TABLET; ORAL

DISOMER

SCHERING

2MG

N011814 001

DISCONTINUED DRUG PRODUCT LIST

6-128(of 430)

** See List Footnote

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

DEXCHLORPHENIRAMINE MALEATE

SYRUP; ORAL			
POLARAMINE			
SCHERING	2MG/5ML	A086837	001 Jul 19, 1982
TABLET; ORAL			
DEXCHLORPHENIRAMINE MALEATE			
ANI PHARMS INC	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
CIPLA	EQ 200MCG BASE/2ML (EQ 100MCG BASE/ML)	A204843	001 Jan 18, 2019

DEXMETHYLPHENIDATE HYDROCHLORIDE

TABLET; ORAL			
DEXMETHYLPHENIDATE HYDROCHLORIDE			
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

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
MYLAN	5MG	A206735	001 Jan 27, 2016
	10MG	A206735	002 Jan 27, 2016
	15MG	A206735	003 Jan 27, 2016
P II	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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

6-129(of 430)

** See List Footnote

DEXTROAMPHETAMINE SULFATE

TABLET; ORAL

DEXEDRINE

GLAXOSMITHKLINE 5MG A084935 001

DEXTROAMPHETAMINE SULFATE

ANI PHARMS INC 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

10MG A040367 001 Oct 31, 2002

PUREPAC PHARM 5MG A084125 001

SANDOZ 10MG A085371 001

VINTAGE PHARMS LLC 5MG A040299 001 May 13, 1999

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 DM

SLATE 15MG/5ML; 6.25MG/5ML A040649 001 Feb 14, 2006

PROMETHAZINE HYDROCHLORIDE AND DEXTROMETHORPHAN HYDROBROMIDE

AMNEAL PHARMS 15MG/5ML; 6.25MG/5ML A090575 001 Feb 08, 2011

+ ANI PHARMS 15MG/5ML; 6.25MG/5ML ** N011265 002 Apr 02, 1984

TRIS PHARMA INC 15MG/5ML; 6.25MG/5ML A091687 001 Jun 28, 2012

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% IN PLASTIC CONTAINER

DHL 5GM/100ML N019971 001 Sep 28, 1995

+ 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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

6-130(of 430)

** See List Footnote

DEXTROSE

INJECTABLE; INJECTION

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/1
00ML; 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/100M
L

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/1
00ML; 140MG/100ML

N019844 001 Jun 10, 1993

ISOLYTE H W/ DEXTROSE 5% IN PLASTIC CONTAINER

B BRAUN 5GM/100ML; 30MG/100ML; 97MG/100ML; 220MG/1
00ML; 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/1
00ML; 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/1
00ML; 530MG/100ML; 500MG/100ML

N018274 001

PLASMA-LYTE 148 AND DEXTROSE 5% IN PLASTIC CONTAINER

+ BAXTER HLTHCARE 5GM/100ML; 30MG/100ML; 37MG/100ML; 368MG/1
00ML; 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

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

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

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 N019870 001 Jun 10, 1993
 /100ML;91MG/100ML

ISOLYTE M W/ DEXTROSE 5% IN PLASTIC CONTAINER
 B BRAUN 5GM/100ML;150MG/100ML;130MG/100ML;280MG N018270 001
 /100ML;91MG/100ML

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 N018840 001 Jun 29, 1983
 /100ML;220MG/100ML

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
 + 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
 5GM/100ML;149MG/100ML;300MG/100ML N018876 006 Mar 28, 1988

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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

DEXTROSE; POTASSIUM CHLORIDE; SODIUM CHLORIDE

INJECTABLE; INJECTION

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
+ ICU MEDICAL INC	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
+ ICU MEDICAL INC	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
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	

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

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
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CARDIOGRAFIN

BRACCO	85%	N011620 002	
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DIATRIZOATE MEGLUMINE

BRACCO	76%	N010040 017	
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HYPAQUE

GE HEALTHCARE	30%	N016403 002	
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	60%	N016403 001	
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RENO-60

BRACCO	60%	N010040 016	
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RENO-DIP

BRACCO	30%	N010040 012	
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UROVIST MEGLUMINE DIU/CT

BAYER HLTHCARE	30%	A087739 001	Sep 23, 1982
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SOLUTION; URETERAL

RENO-30

BRACCO	30%	N010040 021	
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UROVIST CYSTO

BAYER HLTHCARE	30%	A087729 001	Sep 23, 1982
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UROVIST CYSTO PEDIATRIC

BAYER HLTHCARE	30%	A087731 001	Sep 23, 1982
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SOLUTION; URETHRAL

HYPAQUE-CYSTO

GE HEALTHCARE	30%	N016403 003	
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DIATRIZOATE MEGLUMINE; DIATRIZOATE SODIUM

INJECTABLE; INJECTION

ANGIOVIST 292

BAYER HLTHCARE	52%;8%	A087724 001	Sep 23, 1982
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ANGIOVIST 370

BAYER HLTHCARE	66%;10%	A087723 001	Sep 23, 1982
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DIATRIZOATE-60

INTL MEDICATION	52%;8%	A088166 001	Jun 17, 1983
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HYPAQUE-76

GE HEALTHCARE	66%;10%	A086505 001	
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HYPAQUE-M, 75%

GE HEALTHCARE	50%;25%	N010220 003	
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HYPAQUE-M, 90%

GE HEALTHCARE	60%;30%	N010220 002	
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MD-60

MALLINCKRODT	52%;8%	A087074 001	
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MD-76

MALLINCKRODT	66%;10%	A087073 001	
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MD-76R

+ LIEBEL-FLARSHEIM	66%;10%	N019292 001	Sep 29, 1989
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RENOCAL-76

BRACCO	66%;10%	A089347 001	Jun 01, 1988
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RENOGRAFIN-60

BRACCO	52%;8%	N010040 006	
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RENOGRAFIN-76

+ BRACCO	66%;10%	N010040 001	
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Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DIATRIZOATE MEGLUMINE; DIATRIZOATE SODIUM

INJECTABLE; INJECTION

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

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

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

WEST-WARD PHARMS INT 5MG/ML A070311 001 Dec 16, 1985

5MG/ML A070312 001 Dec 16, 1985

5MG/ML A070313 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

DIZAC

PHARMACIA AND UPJOHN 5MG/ML ** N019287 001 Jun 18, 1993

VALIUM

+ ROCHE 5MG/ML ** N016087 001

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

DIAZEPAM

TABLET; ORAL

DIAZEPAM

ACTAVIS ELIZABETH	2MG	A070781 001	Mar 19, 1986
	5MG	A070706 001	Mar 19, 1986
	10MG	A070707 001	Mar 19, 1986
DAVA PHARMS INC	2MG	A070228 002	Sep 26, 1985
	5MG	A070228 003	Sep 26, 1985
	10MG	A070228 001	Sep 26, 1985
DURAMED PHARMS BARR	2MG	A070894 001	Aug 27, 1986
	5MG	A070895 001	Aug 27, 1986
	10MG	A070896 001	Aug 27, 1986
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
NUVO PHARM	10MG	A070464 001	Feb 25, 1986
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
UPSHER SMITH LABS	2MG	A070302 001	Dec 20, 1985
	5MG	A070303 001	Dec 20, 1985
	10MG	A070304 001	Dec 20, 1985
VIRTUS PHARMS	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

DIAZOXIDE

CAPSULE; ORAL

PROGLYCEM

TEVA BRANDED PHARM	50MG	N017425 001	
	100MG	N017425 002	

INJECTABLE; INJECTION

DIAZOXIDE

ABRAXIS PHARM	15MG/ML	A071519 001	Aug 26, 1987
HYPERSTAT			
SCHERING	15MG/ML	N016996 001	

DIBUCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

HEAVY SOLUTION NUPERCAINE

NOVARTIS	2.5MG/ML	N006203 001	
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DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

DICHLORPHENAMIDE

TABLET; ORAL

DARANIDE

+ STRONGBRIDGE US 50MG ** N011366 001

DICLOFENAC EPOLAMINE

SYSTEM; TOPICAL

LICART

+ IBSA INST BIO 1.3% N206976 001 Dec 19, 2018

DICLOFENAC POTASSIUM

TABLET; ORAL

CATAFLAM

+ NOVARTIS 25MG ** N020142 001 Nov 24, 1993

+ 50MG ** N020142 002 Nov 24, 1993

DICLOFENAC POTASSIUM

SANDOZ 50MG A075582 001 Feb 23, 2001

SUN PHARM INDUSTRIES 50MG A075470 001 Feb 21, 2002

WATSON LABS TEVA 50MG A075152 001 Nov 27, 1998

DICLOFENAC SODIUM

SOLUTION; INTRAVENOUS

DICLOFENAC SODIUM

MYLAN LABS LTD 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

RICONPHARMA LLC 1.5% A206715 001 Aug 07, 2017

PENNSAID

+ NUVO PHARMS INC 1.5% ** N020947 001 Nov 04, 2009

SOLUTION/DROPS; OPHTHALMIC

DICLOFENAC SODIUM

APOTEX INC 0.1% A077600 001 Nov 13, 2008

FALCON PHARMS 0.1% N020809 001 May 04, 1998

TABLET, DELAYED RELEASE; ORAL

DICLOFENAC SODIUM

ALLIED 50MG A074986 001 Feb 26, 1999

75MG A074986 002 Feb 26, 1999

CASI PHARMS INC 25MG A074376 001 Sep 28, 1995

50MG A074376 002 Sep 28, 1995

75MG A074394 001 Nov 30, 1995

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

MYLAN 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

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

DICLOXACILLIN SODIUM

CAPSULE; ORAL

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
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DYNAPEN

APOTHECON	EQ 62.5MG BASE/5ML	N050337 002
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PATHOCIL

WYETH AYERST	EQ 62.5MG BASE/5ML	N050092 001
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DICUMAROL

CAPSULE; ORAL

DICUMAROL

LILLY	25MG	N005509 003
	50MG	N005509 001

TABLET; ORAL

DICUMAROL

ABBVIE	25MG	N005545 003
	50MG	N005545 004
	100MG	N005545 005

DICYCLOMINE HYDROCHLORIDE

CAPSULE; ORAL

BENTYL

+ ALLERGAN	10MG	N007409 003 Oct 15, 1984
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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

DICYCLOMINE HYDROCHLORIDE

DR REDDYS	10MG/ML	A080614 001 Feb 11, 1986
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SYRUP; ORAL

BENTYL

+ APTALIS PHARMA US	10MG/5ML **	N007961 002 Oct 15, 1984
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DICYCLOMINE HYDROCHLORIDE

ALPHARMA US PHARMS	10MG/5ML	A084479 001
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TABLET; ORAL

BENTYL

+ ALLERGAN	20MG	N007409 001 Oct 15, 1984
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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

BARR	200MG	A077167 001 Dec 03, 2004
	250MG	A077167 002 Dec 03, 2004
	400MG	A077167 003 Dec 03, 2004
MYLAN PHARMS INC	125MG	A090788 001 Apr 08, 2010
	200MG	A090788 002 Apr 08, 2010
	250MG	A090788 003 Apr 08, 2010
	400MG	A090788 004 Apr 08, 2010

FOR SOLUTION; ORAL

DIDANOSINE

AUROBINDO PHARMA	10MG/ML	A078112 001 Mar 08, 2007
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VIDEX

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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

6-138(of 430)

** See List Footnote

DIRIDANOSINETABLET, 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

DIRIDANOSINE

AUROBINDO	100MG	A077275 001	Aug 14, 2012
	150MG	A077275 002	Aug 14, 2012
	200MG	A077275 003	Aug 14, 2012

DIENESTROL

CREAM;VAGINAL

DIENESTROL

ORTHO MCNEIL PHARM	0.01%	N006110 005	
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DV

SANOFI AVENTIS US	0.01%	A083518 001	
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ESTRAGUARD

SOLVAY	0.01%	A084436 001	
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SUPPOSITORY;VAGINAL

DV

SANOFI AVENTIS US	0.7MG	A083517 001	
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DIETHYLCARBAMAZINE CITRATE

TABLET;ORAL

HETRAZAN

LEDERLE	50MG	N006459 001	
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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

SANOFI AVENTIS US	25MG	N017668 001	
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+

TEVA BRANDED PHARM	25MG	N011722 002	
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TEPANIL

3M	25MG	N011673 001	
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TABLET, EXTENDED RELEASE;ORAL

TENUATE

SANOFI AVENTIS US	75MG	N017669 001	
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TENUATE DOSPAN

+

TEVA BRANDED PHARM	75MG	N012546 001	
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TEPANIL TEN-TAB

3M	75MG	N017956 001	
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DIETHYLSTILBESTROL

INJECTABLE;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	
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	0.5MG	N004040 002	
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STILBESTROL

BRISTOL MYERS SQUIBB	0.1MG	N004056 001	
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	0.5MG	N004056 002	
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TABLET;ORAL

DIETHYLSTILBESTROL

LILLY	0.1MG	N004041 002	
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Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

6-139(of 430)

** See List Footnote

DIETHYLSTILBESTROL

TABLET; ORAL

DIETHYLSTILBESTROL	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
	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			
FOUGERA PHARMS	0.05%	A075187 001	Mar 30, 1998

FLORONE

PHARMACIA AND UPJOHN	0.05% **	N017741 001
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FLORONE E

PHARMACIA AND UPJOHN	0.05%	N019259 001	Aug 28, 1985
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PSORCON

+ TARO PHARMS NORTH	0.05% **	N020205 001	Nov 20, 1992
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OINTMENT; TOPICAL

PSORCON			
+ PHARMACIA AND UPJOHN	0.05%	N019260 001	Aug 28, 1985

PSORCON E

PHARMACIA AND UPJOHN	0.05%	N017994 001
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DIFLUNISAL

TABLET; ORAL

DIFLUNISAL			
ANI PHARMS INC	500MG	A074604 001	Jun 10, 1996
PUREPAC PHARM	250MG	A074285 001	May 07, 1996
	500MG	A074285 002	May 07, 1996
SOCORRO	250MG	A073562 001	Nov 27, 1992
	500MG	A073563 001	Nov 27, 1992
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
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+	500MG **	N018445 002	Apr 19, 1982
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Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

6-140(of 430)

** 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

LANOXIN

+ CONCORDIA

0.1875MG

N020405 003 Sep 30, 1997

0.375MG

N020405 005 Sep 30, 1997

0.5MG

N020405 006 Sep 30, 1997

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

360MG

N020939 005 Sep 14, 2001

420MG

N020939 006 Sep 14, 2001

MYLAN

120MG

A075124 002 Mar 18, 1998

180MG

A075124 003 Mar 18, 1998

240MG

A075124 001 Mar 18, 1998

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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

6-141(of 430)

** See List Footnote

DILTIAZEM HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE;ORAL

DILTIAZEM HYDROCHLORIDE

	420MG	A076563 001	Sep 12, 2006
PAR PHARM	360MG	A209766 001	May 30, 2018
TEVA	60MG	A074079 001	Nov 30, 1993
	90MG	A074079 002	Nov 30, 1993
	120MG	A074079 003	Nov 30, 1993

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
HOSPIRA	5MG/ML	A075004 001	Feb 16, 2000
	5MG/ML	A075106 001	Apr 29, 1999
MYLAN LABS LTD	5MG/ML	A075375 001	Sep 30, 1999

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
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
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
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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
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TABLET; ORAL

DIMENHYDRINATE

HEATHER	50MG	A080841 001
NEXGEN PHARMA INC	50MG	A085985 001
WATSON LABS	50MG	A085166 001

DISCONTINUED DRUG PRODUCT LIST

6-142(of 430)

** See List Footnote

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

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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DIPHENHYDRAMINE HYDROCHLORIDE

CAPSULE; ORAL

DIPHENHYDRAMINE HYDROCHLORIDE

	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	
	50MG	A085083	001	
WHITEWORTH TOWN PLSN	25MG	A083441	001	
	50MG	A080800	001	

ELIXIR; ORAL

BELIX

HALSEY	12.5MG/5ML	A086586	001	Oct 03, 1983
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BENADRYL

MCNEIL CONS	12.5MG/5ML	N005845	004	
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DIBENIL

CENCI	12.5MG/5ML	A088304	001	Dec 16, 1983
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DIPHEN

USL PHARMA	12.5MG/5ML	A084640	001	
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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	
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INJECTABLE; INJECTION

BENADRYL

MCNEIL CONS	10MG/ML	N006146	001	
+	50MG/ML **	N006146	002	

BENADRYL PRESERVATIVE FREE

+	MCNEIL CONS	50MG/ML **	N009486	001
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DIPHENHYDRAMINE HYDROCHLORIDE

BEL MAR	10MG/ML	A080822	001	
DR REDDYS	10MG/ML	A080873	001	
	50MG/ML	A080873	002	
EUROHLTH INTL SARL	50MG/ML	A083183	001	
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
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BELDIN

HALSEY	12.5MG/5ML	A089179	001	Jun 05, 1986
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BENYLIN

PARKE DAVIS	12.5MG/5ML	N006514	004	
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DIPHEN

MORTON GROVE	12.5MG/5ML	A070118	001	Oct 01, 1985
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DIPHENHYDRAMINE HYDROCHLORIDE

ALPHARMA US PHARMS	12.5MG/5ML	A070497	001	Apr 25, 1989
CUMBERLAND SWAN	12.5MG/5ML	A073611	001	Aug 20, 1992
HI TECH PHARMA	12.5MG/5ML	A072416	001	Sep 28, 1990

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

DIPHENHYDRAMINE HYDROCHLORIDE

SYRUP; ORAL

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

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

AKORN 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

HOSPIRA 5MG/ML

A074601 001 Dec 19, 1997

MYLAN LABS LTD 5MG/ML

A075769 001 Nov 27, 2002

IV PERSANTINE

+ BOEHRINGER INGELHEIM 5MG/ML **

N019817 001 Dec 13, 1990

TABLET; ORAL

DIPYRIDAMOLE

ANI PHARMS INC 25MG

A086944 002 Apr 16, 1991

50MG

A086944 001 Feb 25, 1992

75MG

A086944 003 Feb 25, 1992

GLENMARK GENERICS 25MG

A088999 001 Feb 05, 1991

50MG

A089000 001 Feb 05, 1991

75MG

A089001 001 Feb 05, 1991

LANNETT CO INC 25MG

A040898 001 Apr 23, 2008

50MG

A040898 002 Apr 23, 2008

75MG

A040898 003 Apr 23, 2008

OXFORD PHARMS 25MG

A040542 001 Apr 21, 2006

50MG

A040542 002 Apr 21, 2006

75MG

A040542 003 Apr 21, 2006

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

DIRITHROMYCIN

TABLET, DELAYED RELEASE; ORAL

DYNABAC

LILLY RES LABS 250MG

N050678 001 Jun 19, 1995

DISCONTINUED DRUG PRODUCT LIST

6-145(of 430)

** See List Footnote

DISOPYRAMIDE PHOSPHATE

CAPSULE;ORAL

DISOPYRAMIDE PHOSPHATE

AUROLIFE PHARMA LLC	EQ 100MG BASE	A070470 001	Dec 10, 1985
	EQ 150MG BASE	A070471 001	Dec 10, 1985
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
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
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DISULFIRAM

TABLET;ORAL

ANTABUSE

+ TEVA WOMENS	250MG **	N007883 003	
+	500MG **	N007883 002	

DISULFIRAM

HIKMA	250MG	A202652 001	Feb 05, 2014
	500MG	A202652 002	Feb 05, 2014
MYLAN	250MG	A203916 001	Mar 04, 2015
	500MG	A203916 002	Mar 04, 2015
PAR PHARM	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

MYLAN	EQ 125MG VALPROIC ACID	A090407 001	Mar 28, 2011
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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

MYLAN	EQ 125MG VALPROIC ACID	A077254 001	Jul 29, 2008
	EQ 125MG VALPROIC ACID	A090062 001	Mar 17, 2009
	EQ 250MG VALPROIC ACID	A077254 002	Jul 29, 2008
	EQ 250MG VALPROIC ACID	A090062 002	Mar 17, 2009
	EQ 500MG VALPROIC ACID	A077254 003	Jul 29, 2008
	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

ACP NIMBLE	EQ 500MG VALPROIC ACID	A078700 001	Aug 03, 2009
ANCHEN PHARMS	EQ 250MG VALPROIC ACID	A078445 001	Feb 26, 2009
	EQ 500MG VALPROIC ACID	A078445 002	Aug 04, 2009

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 1.25GM BASE/100ML	A074634 001	Sep 27, 1996
LUITPOLD	EQ 12.5MG BASE/ML	A074545 001	Jun 25, 1998
TELIGENT PHARMA INC	EQ 12.5MG BASE/ML	A074098 001	Feb 21, 1995
TEVA PARENTERAL	EQ 12.5MG BASE/ML	A074206 001	Oct 19, 1993

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

DOBUTAMINE HYDROCHLORIDE

INJECTABLE; INJECTION

DOBUTAMINE HYDROCHLORIDE

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 PHARMA GLOBAL	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
CIPLA	20MG/2ML (10MG/ML)	A209634	001	Aug 24, 2018
	80MG/8ML (10MG/ML)	A209634	002	Aug 24, 2018
	160MG/16ML (10MG/ML)	A209634	003	Aug 24, 2018
+ HOSPIRA INC	120MG/6ML (20MG/ML)	N022234	006	Jun 23, 2016
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
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
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DOLASETRON MESYLATE

INJECTABLE; INJECTION

ANZEMET

+ US PHARM HOLDINGS	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

+ US PHARM HOLDINGS	50MG	N020623	001	Sep 11, 1997
+	100MG	N020623	002	Sep 11, 1997

DONEPEZIL HYDROCHLORIDE

SOLUTION; ORAL

ARICEPT

EISAI INC	5MG/5ML	N021719	001	Oct 18, 2004
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TABLET; ORAL

DONEPEZIL HYDROCHLORIDE

ACCORD HLTHCARE	5MG	A201335	001	Aug 29, 2011
	10MG	A201335	002	Aug 29, 2011
APOTEX	5MG	A078841	001	Jun 02, 2011
	10MG	A078841	002	Jun 02, 2011
HIKMA PHARMS	5MG	A090247	001	May 31, 2011
	10MG	A090247	002	May 31, 2011
OSMOTICA PHARM US	23MG	A203114	001	Jan 26, 2016
PAR PHARM	23MG	A202542	001	Jul 24, 2013
SUN PHARM	23MG	A204293	001	Jun 05, 2015
SUN PHARM INDS LTD	5MG	A076786	001	Nov 26, 2010
	10MG	A076786	002	Nov 26, 2010
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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

DONEPEZIL HYDROCHLORIDE

TABLET, ORALLY DISINTEGRATING;ORAL

DONEPEZIL HYDROCHLORIDE

HERITAGE PHARMA	5MG	A078388 002	Nov 26, 2010
	10MG	A078388 001	Nov 26, 2010
SUN PHARM INDUSTRIES	5MG	A077975 002	Dec 11, 2009
	10MG	A077975 001	Dec 11, 2009
ZYDUS PHARMS USA INC	5MG	A090175 001	May 10, 2011
	10MG	A090175 002	May 10, 2011

DONEPEZIL HYDROCHLORIDE; MEMANTINE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE;ORAL

MEMANTINE HYDROCHLORIDE AND DONEPEZIL HYDROCHLORIDE

AMNEAL PHARMS	10MG;14MG	A208328 001	Jan 27, 2017
	10MG;28MG	A208328 002	Jan 27, 2017

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 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
RUBICON	EQ 2% BASE	A078395 001	Oct 28, 2008
TEVA PHARMS	EQ 2% BASE	A078756 001	Dec 04, 2008
WATSON LABS INC	EQ 2% BASE	A202053 001	Sep 11, 2014
ZAMBON SPA	EQ 2% BASE	A091034 001	Dec 04, 2013

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

DORZOLAMIDE HYDROCHLORIDE; TIMOLOL MALEATE

SOLUTION/DROPS;OPHTHALMIC

DORZOLAMIDE HYDROCHLORIDE AND TIMOLOL MALEATE

FDC LTD	EQ 2% BASE;EQ 0.5% BASE	A205295 001	Jun 13, 2019
LANNETT CO INC	EQ 2% BASE;EQ 0.5% BASE	A201998 001	Dec 17, 2014
RUBICON	EQ 2% BASE;EQ 0.5% BASE	A078201 001	Oct 28, 2008
WATSON LABS INC	EQ 2% BASE;EQ 0.5% BASE	A202054 001	Sep 03, 2014
ZAMBON SPA	EQ 2% BASE;EQ 0.5% BASE	A091180 001	Dec 04, 2013

DOXACURIUM CHLORIDE

INJECTABLE;INJECTION

NUROMAX

ABBVIE	EQ 1MG BASE/ML	N019946 001	Mar 07, 1991
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DOXAPRAM HYDROCHLORIDE

INJECTABLE;INJECTION

DOXAPRAM HYDROCHLORIDE

WATSON LABS	20MG/ML	A073529 001	Jan 30, 1992
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DOXAZOSIN MESYLATE

TABLET;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 INC	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
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
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
YAOPHARMA CO LTD	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

DOXEPIIN HYDROCHLORIDE

CAPSULE;ORAL

DOXEPIIN 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
	EQ 150MG BASE	A071676 001	Jan 05, 1988
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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DOXEPIN HYDROCHLORIDE

CAPSULE; ORAL

DOXEPIN HYDROCHLORIDE

	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

SINEQUAN

+ PFIZER

EQ 10MG BASE/ML **

N017516 001

TABLET; ORAL

DOXEPIN HYDROCHLORIDE

MYLAN

EQ 3MG BASE

A202337 001 Jan 20, 2016

EQ 6MG BASE

A202337 002 Jan 20, 2016

DOXERCALCIFEROL

INJECTABLE; INJECTION

DOXERCALCIFEROL

+ HOSPIRA INC

10MCG/5ML (2MCG/ML)

N208614 002 Jul 24, 2018

DOXORUBICIN HYDROCHLORIDE

INJECTABLE; INJECTION

ADRIAMYCIN PFS

PHARMACIA AND UPJOHN

2MG/ML

A063165 001 Jan 30, 1991

200MG/100ML

A063165 002 Jan 30, 1991

DOXORUBICIN HYDROCHLORIDE

ALVOGEN INC

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

2MG/ML

A200901 001 Feb 14, 2012

10MG/VIAL

A200170 001 Oct 28, 2011

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

DOXORUBICIN HYDROCHLORIDE

INJECTABLE; INJECTION

DOXORUBICIN HYDROCHLORIDE

PHARMACIA AND UPJOHN	10MG/VIAL	N050467	001	
	20MG/VIAL	N050467	003	May 20, 1985
	50MG/VIAL	N050467	002	
	150MG/VIAL	N050467	004	Jul 22, 1987
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

MYLAN PHARMS INC	EQ 150MG BASE	A202778	001	Jun 08, 2012
PAR PHARM	EQ 75MG BASE	A065055	004	Apr 18, 2005
SANDOZ INC	EQ 50MG BASE	A065032	001	Jun 30, 2000
	EQ 100MG BASE	A065032	002	Jun 30, 2000
WATSON LABS	EQ 50MG BASE	A065041	001	Apr 28, 2000
	EQ 100MG BASE	A065041	002	Apr 28, 2000
MONODOX				
+ ALMIRALL	EQ 50MG BASE	N050641	002	Feb 10, 1992
+	EQ 75MG BASE	N050641	003	Oct 18, 2006
+	EQ 100MG BASE	N050641	001	Dec 29, 1989

FOR SUSPENSION; ORAL

DOXYCHEL

RACHELLE	EQ 25MG BASE/5ML	A061720	001	
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TABLET; ORAL

DOXYCYCLINE

MYLAN	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
PAR PHARM	EQ 50MG BASE	A065070	001	Dec 15, 2000
	EQ 75MG BASE	A065070	003	Dec 30, 2002
	EQ 100MG BASE	A065070	002	Dec 15, 2000
	EQ 150MG BASE	A065070	004	Jul 14, 2005
SANDOZ INC	EQ 50MG BASE	A065353	001	Nov 27, 2006
	EQ 75MG BASE	A065353	002	Nov 27, 2006
	EQ 100MG BASE	A065353	003	Nov 27, 2006
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 HYCLATE

CAPSULE; ORAL

ACTICLATE CAP

+ ALMIRALL	EQ 75MG BASE	N208253	001	Apr 26, 2016
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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
MYLAN	EQ 50MG BASE	A062337	001	Mar 29, 1982
	EQ 100MG BASE	A062337	002	Mar 29, 1982
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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

DOXYCYCLINE HYCLATE

CAPSULE;ORAL

DOXYCYCLINE HYCLATE

RANBAXY	EQ 50MG BASE	A062479 001	Dec 23, 1983
	EQ 100MG BASE	A062479 002	Dec 23, 1983
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	

PERIOSTAT

+ COLLAGENEX	EQ 20MG BASE **	N050744 001	Sep 30, 1998
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VIBRAMYCIN

+ PFIZER	EQ 50MG BASE **	N050007 001	
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CAPSULE, COATED PELLETS;ORAL

DOXYCYCLINE HYCLATE

PLIVA	EQ 100MG BASE	A063187 001	Jun 30, 1992
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CAPSULE, DELAYED RELEASE;ORAL

DORYX

+ MAYNE PHARMA INTL	EQ 75MG BASE	N050582 002	Aug 13, 2001
+ WARNER CHILCOTT	EQ 100MG BASE	N050582 001	Jul 22, 1985
	EQ 100MG BASE	A062653 001	Oct 30, 1985

DOXYCYCLINE HYCLATE

MEDICIS	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	
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DOXYCYCLINE

WEST-WARD PHARMS INT	EQ 100MG BASE/VIAL	A062450 001	Oct 27, 1983
	EQ 200MG BASE/VIAL	A062450 002	Oct 27, 1983
	EQ 200MG BASE/VIAL	A062569 002	Mar 09, 1988

DOXYCYCLINE HYCLATE

WEST-WARD PHARMS INT	EQ 100MG BASE/VIAL	A062992 001	Feb 16, 1989
	EQ 200MG BASE/VIAL	A062992 002	Feb 16, 1989

VIBRAMYCIN

+ PFIZER	EQ 100MG BASE/VIAL **	N050442 002	
+ PFIZER	EQ 200MG BASE/VIAL **	N050442 001	

TABLET;ORAL

DOXY-LEMMON

TEVA	EQ 100MG BASE	A062581 001	Mar 15, 1985
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DOXYCYCLINE HYCLATE

HEATHER	EQ 100MG BASE	A062462 001	May 11, 1983
INTERPHARM	EQ 100MG BASE	A062764 001	Sep 02, 1988
MUTUAL PHARM	EQ 100MG BASE	A062391 001	Sep 30, 1982
SUPERPHARM	EQ 100MG BASE	A062494 001	Feb 20, 1985
VINTAGE PHARMS	EQ 100MG BASE	A062538 001	Apr 07, 1986
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
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PERIOSTAT

+ GALDERMA LABS LP	EQ 20MG BASE **	N050783 001	Feb 02, 2001
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VIBRA-TABS

+ PFIZER	EQ 100MG BASE **	N050533 001	
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TABLET, DELAYED RELEASE;ORAL

DORYX

+ MAYNE PHARMA	EQ 80MG BASE	N050795 004	Apr 11, 2013
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DORYX MPC

+ MAYNE PHARMA	EQ 60MG BASE **	N050795 007	May 20, 2016
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DOXYCYCLINE HYCLATE

IMPAX LABS INC	EQ 75MG BASE	A090505 001	Dec 28, 2010
	EQ 100MG BASE	A090505 002	Dec 28, 2010
MYLAN	EQ 75MG BASE	A090431 001	Dec 28, 2010
	EQ 80MG BASE	A090431 004	Apr 29, 2016

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

DOXYCYCLINE HYCLATETABLET, DELAYED RELEASE;ORAL
DOXYCYCLINE HYCLATE

	EQ 100MG BASE	A090431 002	Dec 28, 2010
	EQ 200MG BASE	A090431 005	May 19, 2016
MYLAN PHARMS INC	EQ 150MG BASE	A091052 001	Feb 08, 2012
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 SUCCINATECAPSULE;ORAL
UNISOM

PFIZER	25MG	N019440 001	Feb 05, 1986
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TABLET;ORAL

DECAPRYN

SANOFI AVENTIS US	12.5MG	N006412 015	
	25MG	N006412 014	

DOXY-SLEEP-AID

PAR PHARM	25MG	A070156 001	Jul 02, 1987
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DOXYLAMINE SUCCINATE

COPLEY 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	
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DROMOSTANOLONE PROPIONATE

INJECTABLE;INJECTION

DROLBAN

LILLY	50MG/ML	N012936 001	
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DRONABINOL

CAPSULE;ORAL

DRONABINOL

INSYS THERAP	2.5MG	A078501 001	Aug 19, 2011
	5MG	A078501 002	Aug 19, 2011
	10MG	A078501 003	Aug 19, 2011

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	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
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

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			
AKORN MFG	2.5MG/ML;EQ 0.05MG BASE/ML	N016049 001	

DISCONTINUED DRUG PRODUCT LIST

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** 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

DUTASTERIDE

CAPSULE;ORAL

DUTASTERIDE

ACTAVIS LABS FL INC	0.5MG	A202808 001	Nov 20, 2015
APOTEX INC	0.5MG	A204292 001	Nov 24, 2015
BRECKENRIDGE	0.5MG	A204705 001	Nov 20, 2015
HIKMA	0.5MG	A202204 001	Nov 23, 2015
MYLAN	0.5MG	A203241 001	Jun 14, 2016
PII	0.5MG	A208227 001	Jun 22, 2018
RISING	0.5MG	A202530 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	
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INJECTABLE;INJECTION

NEOTHYLLINE

TEVA	250MG/ML	N009088 001	
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TABLET;ORAL

DILOR

SAVAGE LABS	200MG	A084514 001	
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DILOR-400

SAVAGE LABS	400MG	A084751 001	
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LUFYLLIN

MYLAN SPECIALITY LP	200MG	A084566 001	
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	400MG	A084566 002	
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NEOTHYLLINE

TEVA	200MG	N007794 001	
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	400MG	N007794 002	
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ECHOTHIOPHATE IODIDE

FOR SOLUTION;OPHTHALMIC

PHOSPHOLINE IODIDE

WYETH PHARMS	0.03%	N011963 002	
	0.06%	N011963 004	
	0.25%	N011963 003	

ECONAZOLE NITRATE

CREAM;TOPICAL

ECONAZOLE NITRATE

CASI PHARMS INC	1%	A076075 001	Nov 26, 2002
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EDETATE CALCIUM DISODIUM

TABLET;ORAL

CALCIUM DISODIUM VERSENATE

MEDICIS	500MG	N008922 002	
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EDROPHONIUM CHLORIDE

INJECTABLE;INJECTION

EDROPHONIUM CHLORIDE

HOSPIRA	10MG/ML	A040131 001	Feb 24, 1998
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WATSON LABS	10MG/ML	A040044 001	Mar 20, 1996
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EDROPHONIUM CHLORIDE PRESERVATIVE FREE

WATSON LABS	10MG/ML	A040043 001	Mar 20, 1996
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Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

EDROPHONIUM CHLORIDE

INJECTABLE; INJECTION

ENLON

MYLAN INSTITUTIONAL	10MG/ML	A088873	001	Aug 06, 1985
REVERSOL				
ORGANON USA INC	10MG/ML	A089624	001	May 13, 1988
TENSILON				
+ TELIGENT	10MG/ML **	N007959	001	
TENSILON PRESERVATIVE FREE				
+ TELIGENT	10MG/ML **	N007959	002	

EFAVIRENZ

CAPSULE; ORAL

SUSTIVA

+ BRISTOL MYERS SQUIBB	100MG **	N020972	002	Sep 17, 1998
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TABLET; ORAL

SUSTIVA

+ BRISTOL MYERS SQUIBB	300MG **	N021360	001	Feb 01, 2002
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EFAVIRENZ; EMTRICITABINE; TENOFOVIR DISOPROXIL FUMARATE

TABLET; ORAL

EFAVIRENZ, EMTRICITABINE, AND TENOFOVIR DISOPROXIL FUMARATE

TEVA PHARMS USA	600MG; 200MG; 300MG	A091215	001	Nov 09, 2018
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EFAVIRENZ; LAMIVUDINE; TENOFOVIR DISOPROXIL FUMARATE

TABLET; ORAL

EFAVIRENZ, LAMIVUDINE AND TENOFOVIR DISOPROXIL FUMARATE

AUROBINDO PHARMA LTD	600MG; 300MG; 300MG	N022343	001	Aug 15, 2018
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MACLEODS PHARMS LTD	400MG; 300MG; 300MG	N210649	001	Mar 15, 2019
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EFLORNITHINE HYDROCHLORIDE

INJECTABLE; INJECTION

ORNIDYL

SANOFI AVENTIS US	200MG/ML	N019879	002	Nov 28, 1990
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ELTROMBOPAG OLAMINE

TABLET; ORAL

PROMACTA

+ NOVARTIS	EQ 100MG ACID	N022291	005	Nov 16, 2012
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ELVITEGRAVIR

TABLET; ORAL

VITEKTA

+ GILEAD SCIENCES INC	85MG	N203093	001	Sep 24, 2014
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+ GILEAD SCIENCES INC	150MG	N203093	002	Sep 24, 2014
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EMEDASTINE DIFUMARATE

SOLUTION/DROPS; OPHTHALMIC

EMADINE

+ NOVARTIS	0.05%	N020706	001	Dec 29, 1997
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EMTRICITABINE; TENOFOVIR DISOPROXIL FUMARATE

TABLET; ORAL

EMTRICITABINE AND TENOFOVIR DISOPROXIL FUMARATE

AMNEAL PHARMS CO	100MG; 150MG	A209721	001	Aug 22, 2018
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	133MG; 200MG	A209721	002	Aug 22, 2018
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	167MG; 250MG	A209721	003	Aug 22, 2018
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	200MG; 300MG	A209721	004	Aug 22, 2018
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LAURUS LABS LTD	200MG; 300MG	A212114	001	Jul 26, 2019
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MYLAN	200MG; 300MG	A206436	001	Apr 09, 2018
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ENALAPRIL MALEATE

FOR SOLUTION; ORAL

EPANED KIT

+ SILVERGATE PHARMS	1MG/ML	N204308	001	Aug 13, 2013
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TABLET; ORAL

ENALAPRIL MALEATE

APOTHECON	2.5MG	A075583	001	Aug 22, 2000
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	5MG	A075583	002	Aug 22, 2000
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	10MG	A075583	003	Aug 22, 2000
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	20MG	A075583	004	Aug 22, 2000
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BEXIMCO PHARMS USA	2.5MG	A075621	001	Aug 22, 2000
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	5MG	A075621	002	Aug 22, 2000
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	10MG	A075621	003	Aug 22, 2000
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Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

ENALAPRIL MALEATE

TABLET; ORAL

ENALAPRIL MALEATE

	20MG	A075621 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
	2.5MG	A075480 001	Aug 22, 2000
	5MG	A075472 002	Aug 22, 2000
	5MG	A075480 002	Aug 22, 2000
	10MG	A075472 003	Aug 22, 2000
	10MG	A075480 003	Aug 22, 2000
	20MG	A075472 004	Aug 22, 2000
	20MG	A075480 004	Aug 22, 2000
SANDOZ	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
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

IVAX SUB TEVA PHARMS	5MG; 12.5MG	A075736 001	Mar 25, 2003
	10MG; 25MG	A075736 002	Mar 25, 2003
UPSHER SMITH LABS	5MG; 12.5MG	A076116 001	Sep 19, 2001
	10MG; 25MG	A076116 002	Sep 19, 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
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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	

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

ENOXACINTABLET; 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
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ENTACAPONE

TABLET; ORAL

ENTACAPONE

MYLAN PHARMS INC	200MG	A202394 001	May 13, 2013
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ENTECAVIR

TABLET; ORAL

ENTECAVIR

MYLAN	0.5MG	A206226 001	Mar 26, 2019
	1MG	A206226 002	Mar 26, 2019
PAR PHARM INC	0.5MG	A206294 001	Nov 23, 2016
	1MG	A206294 002	Nov 23, 2016

EPINASTINE HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

EPINASTINE HYDROCHLORIDE

CASI PHARMS INC	0.05%	A203384 001	Dec 07, 2016
SUN PHARM INDS	0.05%	A091626 001	Oct 31, 2011

EPINEPHRINE

AEROSOL, METERED; INHALATION

BRONKAID MIST

STERLING	0.25MG/INH	N016803 001	
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EPINEPHRINE

ARMSTRONG PHARMS	0.2MG/INH	A087907 001	May 23, 1984
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PRIMATENE MIST

WYETH CONS	0.2MG/INH	N016126 001	
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INJECTABLE; INJECTION

SUS-PHRINE SULFITE FREE

FOREST LABS	1.5MG/AMP	N007942 003	Feb 05, 1999
	5MG/ML	N007942 001	

INJECTABLE; INTRAMUSCULAR

EPI E Z PEN JR

MYLAN SPECIALITY LP	0.15MG/DELIVERY	N019430 004	Aug 03, 1995
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EPIPEN E Z PEN

MYLAN SPECIALITY LP	0.3MG/DELIVERY	N019430 003	Aug 03, 1995
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INJECTABLE; INTRAMUSCULAR, SUBCUTANEOUS

TWINJECT 0.15

IMPAX	EQ 0.15MG/DELIVERY	N020800 002	May 28, 2004
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TWINJECT 0.3

IMPAX	EQ 0.3MG/DELIVERY	N020800 001	May 30, 2003
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SOLUTION; INTRAMUSCULAR, SUBCUTANEOUS

EPINEPHRINE

AM REGENT	EQ 1MG BASE/ML (EQ 1MG BASE/ML)	A207568 001	Jul 06, 2018
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EPINEPHRINE BITARTRATE

AEROSOL, METERED; INHALATION

BRONITIN MIST

WYETH CONS	0.3MG/INH	N016126 002	
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MEDIHALER-EPI

3M	0.3MG/INH	N010374 003	
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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	

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

EPINEPHRINE BITARTRATE; PRILUCAINE 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

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

ABBOTT 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

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

EPIRUBICIN HYDROCHLORIDE

INJECTABLE; INJECTION

EPIRUBICIN HYDROCHLORIDE

ACTAVIS TOTOWA	10MG/5ML (2MG/ML)	A065445	001	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	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
MYLAN INSTITUTIONAL	50MG/25ML (2MG/ML)	A065371	001	Nov 28, 2007
	200MG/100ML (2MG/ML)	A065371	002	Nov 28, 2007
MYLAN LABS LTD	50MG/25ML (2MG/ML)	A091599	001	Mar 12, 2012
	200MG/100ML (2MG/ML)	A091599	002	Mar 12, 2012
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

INSPIRA

GD SEARLE LLC	100MG	N021437	003	Sep 27, 2002
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EPROSARTAN MESYLATE

TABLET; ORAL

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

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

ERGOCALCIFEROL

CAPSULE; ORAL

DELTALIN

LILLY	50,000 IU	A080884	001	
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VITAMIN D

CHASE CHEM	50,000 IU	A080747	001	
EVERYLIFE	50,000 IU	A080956	001	
IMPAX LABS	50,000 IU	A080951	001	
LANNETT	50,000 IU	A080825	001	
VITARINE	50,000 IU	A084053	001	
WEST WARD	50,000 IU	A083102	001	

ERGOLOID MESYLATES

CAPSULE; ORAL

HYDERGINE LC

NOVARTIS	1MG	N018706	001	Jan 18, 1983
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SOLUTION; ORAL

HYDERGINE

NOVARTIS	1MG/ML	N018418	001	
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TABLET; ORAL

ERGOLOID MESYLATES

MUTUAL PHARM	1MG	A088891	001	Nov 01, 1985
WATSON LABS	1MG	A086433	001	May 27, 1982

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

ERGOLOID MESYLATES

TABLET; ORAL

ERGOLOID MESYLATES	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

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			
MYLAN	2%	A065009 001	Mar 18, 2002

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

ERYTHROMYCIN

GEL;TOPICAL			
EMGEL			
ALTANA	2%	A063107 001	Aug 23, 1991
LOTION;TOPICAL			
E-SOLVE 2			
SYOSSET	2%	A062467 001	Jul 03, 1985
OINTMENT;OPHTHALMIC			
ERYTHROMYCIN			
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	
ERYMAX			
MERZ PHARMS	2%	A062508 002	Jul 11, 1985
ERYTHRA-DERM			
MLV	2%	A062687 001	Feb 05, 1988
ERYTHRO-STATIN			
HI TECH PHARMA	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 AND LOMB	2%	A064039 001	Jan 27, 1994
FOUGERA PHARMS	2%	A064187 001	Sep 30, 1997
LILLY	2%	N050532 001	
PHARMAFAIR	1.5%	A062485 001	Jul 11, 1984
	2%	A062616 001	Jul 25, 1985
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
MYLAN	2%	A064128 001	Jul 03, 1996
T-STAT			
WESTWOOD SQUIBB	2%	A062748 001	Jul 23, 1987
TABLET, COATED PARTICLES;ORAL			
PCE			
+ ARBOR PHARMS LLC	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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

ERYTHROMYCIN

TABLET, DELAYED RELEASE;ORAL

E-MYCIN

ARBOR PHARMS INC	250MG	A060272	001
	333MG	A060272	002

ILOTYCIN

DISTA	250MG	A061910	001
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R-P MYCIN

SOLVAY	250MG	A061659	001
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ROBIMYCIN

ROBINS AH	250MG	A061633	001
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ERYTHROMYCIN ESTOLATE

CAPSULE;ORAL

ERYTHROMYCIN ESTOLATE

BARR	EQ 125MG BASE	A062162	001
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	EQ 250MG BASE	A062162	002
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IVAX SUB TEVA PHARMS	EQ 250MG BASE	A062237	001
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WATSON LABS	EQ 250MG BASE	A062087	001
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ILOSONE

LILLY	EQ 125MG BASE	A061897	001
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	EQ 250MG BASE	A061897	002
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FOR SUSPENSION;ORAL

ILOSONE

DISTA	EQ 125MG BASE/5ML	A061893	001
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SUSPENSION;ORAL

ERYTHROMYCIN ESTOLATE

ACP NIMBLE	EQ 125MG BASE/5ML	A062169	001	Oct 17, 1990
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	EQ 250MG BASE/5ML	A062169	002	Oct 17, 1990
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ALPHARMA US PHARMS	EQ 125MG BASE/5ML	A062353	001	Nov 18, 1982
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	EQ 250MG BASE/5ML	A062409	001	Dec 16, 1982
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LIFE LABS	EQ 250MG BASE/5ML	A062362	001	Dec 17, 1982
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ILOSONE

LILLY	EQ 125MG BASE/5ML	A061894	001
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	EQ 125MG BASE/5ML	N050010	001
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	EQ 250MG BASE/5ML	A061894	002
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	EQ 250MG BASE/5ML	N050010	002
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SUSPENSION/DROPS;ORAL

ILOSONE

LILLY	EQ 100MG BASE/ML	A061894	003
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TABLET;ORAL

ILOSONE

LILLY	EQ 500MG BASE	A061896	001
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TABLET, CHEWABLE;ORAL

ILOSONE

DISTA	EQ 125MG BASE	A061895	001
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	EQ 250MG BASE	A061895	002
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ERYTHROMYCIN ESTOLATE; SULFISOXAZOLE ACETYL

SUSPENSION;ORAL

ILOSONE SULFA

LILLY	EQ 125MG BASE/5ML;EQ 600MG BASE/5ML	N050599	001	Sep 29, 1989
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ERYTHROMYCIN ETHYLSUCCINATE

GRANULE;ORAL

PEDIAMYCIN

ROSS LABS	EQ 200MG BASE/5ML	A062305	001
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SUSPENSION;ORAL

E-MYCIN E

PHARMACIA AND UPJOHN	EQ 200MG BASE/5ML	A062198	001
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	EQ 400MG BASE/5ML	A062198	002
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E.E.S. 200

ARBOR PHARMS LLC	EQ 200MG BASE/5ML **	A061639	001
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E.E.S. 400

ARBOR PHARMS LLC	EQ 400MG BASE/5ML **	A061639	002
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ERYTHROMYCIN ETHYLSUCCINATE

ALPHARMA US PHARMS	EQ 200MG BASE/5ML	A062200	001
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	EQ 400MG BASE/5ML	A062200	002
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DISTA	EQ 200MG BASE/5ML	A062177	001
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	EQ 400MG BASE/5ML	A062177	002
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NASKA	EQ 400MG BASE/5ML	A062674	001	Mar 10, 1987
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Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

ERYTHROMYCIN ETHYLSUCCINATE

SUSPENSION; ORAL

ERYTHROMYCIN ETHYLSUCCINATE

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	
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PEDIAMYCIN 400

ARBOR PHARMS LLC	EQ 400MG BASE/5ML	A062304 002	
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WYAMYCIN E

WYETH AYERST	EQ 200MG BASE/5ML	A062123 002	
	EQ 400MG BASE/5ML	A062123 001	

SUSPENSION/DROPS; ORAL

PEDIAMYCIN

ROSS LABS	EQ 100MG BASE/2.5ML	A062305 002	
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TABLET; ORAL

E.E.S. 400

ARBOR PHARMS LLC	EQ 400MG BASE	A061905 001	
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ERYTHROMYCIN ETHYLSUCCINATE

BARR	EQ 400MG BASE	A062256 001	
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MYLAN	EQ 400MG BASE	A062847 001	Sep 14, 1988
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TABLET, CHEWABLE; ORAL

E.E.S.

ARBOR PHARMS INC	EQ 200MG BASE	N050297 002	
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ERYPED

ARBOR PHARMS INC	EQ 200MG BASE	N050297 003	Jul 05, 1988
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PEDIAMYCIN

ROSS LABS	EQ 200MG BASE	A062306 001	
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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
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ERYZOLE

ALRA	EQ 200MG BASE/5ML; EQ 600MG BASE/5ML	A062758 001	Jun 15, 1988
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PEDIAZOLE

ROSS LABS	EQ 200MG BASE/5ML; EQ 600MG BASE/5ML	N050529 001	
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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
TEVA PARENTERAL	EQ 500MG BASE/VIAL	A063253 001	Jul 30, 1993
	EQ 1GM BASE/VIAL	A063253 002	Jul 30, 1993

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

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

ARBOR PHARMS LLC	EQ 125MG BASE	A060359 002	
	EQ 500MG BASE	A060359 003	

ERYTHROMYCIN STEARATE

ANI PHARMS INC	EQ 250MG BASE	A061461 001	
	EQ 250MG BASE	A061591 001	
	EQ 500MG BASE	A061461 002	
	EQ 500MG BASE	A063179 001	May 15, 1990

LEDERLE	EQ 250MG BASE	A062089 001	
	EQ 500MG BASE	A062089 002	

MYLAN	EQ 250MG BASE	A061505 001	
	EQ 500MG BASE	A061505 002	

PUREPAC PHARM	EQ 250MG BASE	A061743 001	
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WATSON LABS	EQ 250MG BASE	A062121 002	
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	EQ 500MG BASE	A062121 001	
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ETHRIL 250

BRISTOL MYERS SQUIBB	EQ 250MG BASE	A061605 001	
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ETHRIL 500

BRISTOL MYERS SQUIBB	EQ 500MG BASE	A061605 002	
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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

MYLAN PHARMS INC	EQ 5MG BASE	A077660 001	Jul 31, 2007
	EQ 10MG BASE	A077660 002	Jul 31, 2007
	EQ 20MG BASE	A077660 003	Jul 31, 2007

TABLET;ORAL

ESCITALOPRAM OXALATE

MYLAN	EQ 5MG BASE	A077550 001	May 14, 2015
	EQ 10MG BASE	A077550 002	May 14, 2015
	EQ 20MG BASE	A077550 003	May 14, 2015

ESMOLOL HYDROCHLORIDE

INJECTABLE;INJECTION

BREVIBLOC

BAXTER HLTHCARE	10MG/ML	N019386 003	Aug 15, 1988
	20MG/ML	N019386 007	May 28, 2003

ESMOLOL HYDROCHLORIDE

AM REGENT	10MG/ML	A201126 001	Feb 20, 2015
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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
HEC PHARM	EQ 20MG BASE	A207265 002	May 18, 2018
	EQ 40MG BASE	A207265 001	May 18, 2018

TABLET, DELAYED RELEASE;ORAL

ESOMEPRAZOLE MAGNESIUM

P AND L	EQ 20MG BASE	A209202 001	Mar 05, 2019
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DISCONTINUED DRUG PRODUCT LIST

6-164(of 430)

** See List Footnote

ESOMEPRAZOLE SODIUM

INJECTABLE; INTRAVENOUS

ESOMEPRAZOLE SODIUM

AUROBINDO PHARMA LTD	EQ 20MG BASE/VIAL	A204657	001	Aug 10, 2016
MYLAN LABS LTD	EQ 20MG BASE/VIAL	A202686	001	May 17, 2017
SUN PHARMA GLOBAL	EQ 20MG BASE/VIAL	A200882	001	Mar 18, 2013
NEXIUM IV				
+ ASTRAZENECA PHARMS	EQ 20MG BASE/VIAL **	N021689	001	Mar 31, 2005

ESOMEPRAZOLE STRONTIUM

CAPSULE, DELAYED RELEASE; ORAL

ESOMEPRAZOLE STRONTIUM

+ R2 PHARMA LLC	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

ESTRADIOL

FILM, EXTENDED RELEASE; 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

FEMPATCH

PARKE DAVIS	0.025MG/24HR	N020417	001	Dec 03, 1996
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VIVELLE

NOVARTIS	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

GEL; TOPICAL

ESTROGEL

ASCEND THERAPS US	0.06%	N021166	001	Feb 09, 2004
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TABLET; ORAL

ESTRACE

BRISTOL MYERS SQUIBB	0.5MG	A081295	001	Jun 30, 1993
	1MG	A084499	001	
	2MG	A084500	001	

ESTRADIOL

LANNETT HOLDINGS INC	0.5MG	A040138	001	Jan 30, 1998
	1MG	A040138	002	Jan 30, 1998
	2MG	A040138	003	Jan 30, 1998
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

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

ESTRADIOLTABLET; VAGINAL
VAGIFEM

+ NOVO NORDISK INC 25MCG ** N020908 001 Mar 26, 1999

ESTRADIOL ACETATETABLET; ORAL
FEMTRACE+ APIL 0.45MG N021633 001 Aug 20, 2004
+ 0.9MG N021633 002 Aug 20, 2004
+ 1.8MG N021633 003 Aug 20, 2004ESTRADIOL CYPIONATEINJECTABLE; INJECTION
DEPO-ESTRADIOLPHARMACIA AND UPJOHN 1MG/ML A085470 001
3MG/ML A085470 002

ESTRADIOL CYPIONATE

DR REDDYS 5MG/ML A085620 001

ESTRADIOL CYPIONATE; MEDROXYPROGESTERONE ACETATEINJECTABLE; INTRAMUSCULAR
LUNELLE

PHARMACIA AND UPJOHN 5MG/0.5ML; 25MG/0.5ML N020874 001 Oct 05, 2000

ESTRADIOL CYPIONATE; TESTOSTERONE CYPIONATEINJECTABLE; 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 HEMIHYDRATEEMULSION; TOPICAL
ESTRASORB

+ EXELTIS USA INC 0.25% N021371 001 Oct 09, 2003

ESTRADIOL VALERATEINJECTABLE; INJECTION
ESTRADIOL VALERATEDR 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 001ESTRADIOL VALERATE; TESTOSTERONE ENANTHATEINJECTABLE; 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 001ESTRADIOL; NORETHINDRONE ACETATE

TABLET; ORAL

ESTRADIOL AND NORETHINDRONE ACETATE

ACCORD HLTHCARE 1MG; 0.5MG A210233 001 Feb 28, 2018

TEVA PHARMS USA 0.5MG; 0.1MG A200747 001 Mar 08, 2012

ESTRADIOL; NORGESTIMATE

TABLET; ORAL

PREFEST

+ TEVA WOMENS 1MG, 1MG; N/A, 0.09MG ** N021040 001 Oct 22, 1999

ESTROGENS, CONJUGATED

TABLET; ORAL

PREMARIN

WYETH PHARMS 2.5MG N004782 002

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

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
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PREMPRO (PREMARIN;CYCRIN)

WYETH PHARMS INC	0.625MG,0.625MG;2.5MG,2.5MG	N020303 001	Dec 30, 1994
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ESTROGENS, CONJUGATED; MEPROBAMATE

TABLET;ORAL

MILPREM-200

MEDPOINTE PHARM HLC	0.45MG;200MG	N011045 002
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MILPREM-400

MEDPOINTE PHARM HLC	0.45MG;400MG	N011045 001
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PMB 200

WYETH AYERST	0.45MG;200MG	N010971 005
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PMB 400

WYETH AYERST	0.45MG;400MG	N010971 003
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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

ESTRONE

INJECTABLE;INJECTION

ESTROGENIC SUBSTANCE

WYETH AYERST	2MG/ML	A083488 001
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ESTRONE

DR REDDYS	5MG/ML	A085239 001
WATSON LABS	2MG/ML	A083397 001

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

ESTRONE

INJECTABLE; INJECTION

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

MYLAN 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

PHARMACIA AND UPJOHN 0.75MG

A083220 001

OGEN 1.25

PHARMACIA AND UPJOHN 1.5MG

A083220 002

OGEN 2.5

PHARMACIA AND UPJOHN 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

BRECKENRIDGE 1MG

A203087 001 May 08, 2019

2MG

A203087 002 May 08, 2019

3MG

A203087 003 May 08, 2019

HIKMA 1MG

A091153 001 Apr 15, 2014

2MG

A091153 002 Apr 15, 2014

3MG

A091153 003 Apr 15, 2014

WOCKHARDT LTD 1MG

A091165 001 Jul 14, 2011

2MG

A091165 002 Jul 14, 2011

3MG

A091165 003 Jul 14, 2011

ETHACRYNIC ACID

TABLET; ORAL

EDECIN

BAUSCH 50MG

N016092 002

ETHAMBUTOL HYDROCHLORIDE

TABLET; ORAL

MYAMBUTOL

STI PHARMA LLC 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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

ETHCHLORVYNOLCAPSULE;ORAL
PLACIDYL

500MG	N010021 002
750MG	N010021 010

ETHINAMATECAPSULE;ORAL
VALMID

DISTA	500MG	N009750 001
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ETHINYL ESTRADIOLTABLET;ORAL
ESTINYL

SCHERING	0.02MG	N005292 001
	0.05MG	N005292 002
	0.5MG	N005292 003

FEMINONE

PHARMACIA AND UPJOHN	0.05MG	N016649 001
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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
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DEMULEN 1/50-21

GD SEARLE LLC	0.05MG;1MG	N016927 001
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ZOVIA 1/35E-21

WATSON PHARMS TEVA	0.035MG;1MG	A072720 001	Dec 30, 1991
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ZOVIA 1/50E-21

WATSON LABS	0.05MG;1MG	A072722 001	Dec 30, 1991
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TABLET;ORAL-28

DEMULEN 1/35-28

GD SEARLE LLC	0.035MG;1MG **	N018160 001
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DEMULEN 1/50-28

GD SEARLE LLC	0.05MG;1MG **	N016936 001
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ETHINYL ESTRADIOL; FERROUS FUMARATE; NORETHINDRONE

TABLET;ORAL-28

NORQUEST FE

GD SEARLE LLC	0.035MG;75MG;1MG	N018926 001	Jul 18, 1986
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ETHINYL ESTRADIOL; FERROUS FUMARATE; NORETHINDRONE ACETATE

TABLET;ORAL-28

NORLESTRIN FE 1/50

PARKE DAVIS	0.05MG;75MG;1MG	N016766 001
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NORLESTRIN FE 2.5/50

PARKE DAVIS	0.05MG;75MG;2.5MG	N016854 001
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ETHINYL ESTRADIOL; LEVONORGESTREL

TABLET;ORAL

LYBREL

+ WYETH PHARMS INC	0.02MG;0.09MG **	N021864 001	May 22, 2007
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PREVEN EMERGENCY CONTRACEPTIVE KIT

TEVA BRANDED PHARM	0.05MG;0.25MG	N020946 001	Sep 01, 1998
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TABLET;ORAL-21

ALESSE

+ CADENCE HEALTH	0.02MG;0.1MG **	N020683 001	Mar 27, 1997
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AVIANE-21

DURAMED PHARMS BARR	0.02MG;0.1MG	A075796 002	Apr 30, 2001
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ENPRESSE-21

DURAMED PHARMS BARR	0.03MG,0.04MG,0.03MG;0.05MG,0.075MG,0.125MG	A075809 001	Jul 16, 2001
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LESSINA-21

BARR	0.02MG;0.1MG	A075803 001	Mar 20, 2002
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LEVLITE

+ BAYER HLTHCARE	0.02MG;0.1MG **	N020860 001	Jul 13, 1998
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LEVONORGESTREL AND ETHINYL ESTRADIOL

BARR	0.02MG;0.1MG	A075862 001	Apr 29, 2003
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Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

ETHINYL ESTRADIOL; LEVONORGESTREL

TABLET; ORAL-21

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					
MAYNE PHARMA	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
LEVLITE					
+ BAYER HLTHCARE	0.02MG;0.1MG **		N020860	002	Jul 13, 1998
LEVONORGESTREL AND ETHINYL ESTRADIOL					
BARR	0.02MG;0.1MG		A075862	002	Apr 29, 2003
MYLAN LABS LTD	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
TRIPHASIL-28					
+ WYETH PHARMS INC	0.03MG,0.04MG,0.03MG;0.05MG,0.075MG,0.125MG **		N019190	001	Nov 01, 1984

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-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
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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

ETHINYL ESTRADIOL; NORETHINDRONE

TABLET; ORAL-21			
OVCON-35			
+ WARNER CHILCOTT	0.035MG; 0.4MG **	N018127	001
OVCON-50			
WARNER CHILCOTT	0.05MG; 1MG	N018128	001
TRI-NORINYL 21-DAY			
MAYNE PHARMA	0.035MG, 0.035MG, 0.035MG; 0.5MG, 1MG, 0.5MG	N018977	001 Apr 13, 1984
TABLET; ORAL-28			
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
NORETHIN 1/35E-28			
WATSON LABS	0.035MG; 1MG	A071481	001 Apr 12, 1988
NORETHINDRONE AND ETHINYL ESTRADIOL			
MYLAN LABS LTD	0.035MG; 0.4MG	A200897	001 May 11, 2015
	0.035MG, 0.035MG, 0.035MG; 0.5MG, 0.75MG, 1MG	A200486	001 Dec 28, 2015
	0.035MG; 0.5MG	A200488	001 Oct 21, 2015
	0.035MG; 1MG	A200489	001 Oct 21, 2015
	0.05MG; 1MG	A203006	001 Aug 05, 2013
WATSON LABS	0.035MG, 0.035MG, 0.035MG; 0.5MG, 0.75MG, 1MG	A076393	001 Feb 04, 2010
NORETHINDRONE AND ETHINYL ESTRADIOL (7/14)			
WATSON LABS	0.035MG, 0.035MG; 0.5MG, 1MG	A071042	001 Sep 24, 1991
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
OVCON-35			
+ WARNER CHILCOTT LLC	0.035MG; 0.4MG **	N017716	001
OVCON-50			
WARNER CHILCOTT LLC	0.05MG; 1MG **	N017576	001
TABLET, CHEWABLE; ORAL			
FEMCON FE			
+ APIL	0.035MG; 0.4MG	N021490	001 Nov 14, 2003

ETHINYL ESTRADIOL; NORETHINDRONE ACETATE

TABLET; ORAL			
FEMHRT			
+ APIL	0.005MG; 1MG **	N021065	002 Oct 15, 1999
NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL AND ETHINYL ESTRADIOL AND FERROUS FUMARATE			
MYLAN LABS LTD	0.01MG, 0.01MG; 1MG, N/A	A205049	001 May 31, 2016
NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL AND FERROUS FUMARATE			
APOTEX	0.02MG; 1MG	A208639	001 Mar 21, 2018
TABLET; ORAL-21			
ESTROSTEP 21			
+ APIL	0.02MG, 0.03MG, 0.035MG; 1MG, 1MG, 1MG **	N020130	001 Oct 09, 1996
NORLESTRIN 21 1/50			
PARKE DAVIS	0.05MG; 1MG	N016749	001
NORLESTRIN 21 2.5/50			
PARKE DAVIS	0.05MG; 2.5MG	N016852	001
TABLET; ORAL-28			
NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL			
MYLAN LABS LTD	0.02MG, 0.03MG, 0.035MG; 1MG, 1MG, 1MG	A205069	001 Jun 22, 2018
NORLESTRIN 28 1/50			
PARKE DAVIS	0.05MG; 1MG	N016723	001
TABLET, CHEWABLE; ORAL			
NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL AND FERROUS FUMARATE			
MYLAN LABS LTD	0.02MG; 1MG	A206120	001 Sep 12, 2017
TABLET, CHEWABLE, TABLET; ORAL			
LO MINASTRIN FE			
+ APIL	0.01MG, 0.01MG, N/A; 1MG, N/A, N/A	N204654	001 Jul 24, 2013

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

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

MYLAN LABS LTD 0.025MG,0.025MG,0.025MG;0.18MG,0.215MG,
0.25MG A202132 001 Sep 09, 2015WATSON 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, A076626 001 Aug 17, 2006

0.25MG

0.035MG;0.25MG A076627 001 Aug 17, 2006

ORTHO CYCLEN-28

+ JANSSEN PHARMS 0.035MG;0.25MG N019653 002 Dec 29, 1989

ETHINYL ESTRADIOL; NORGESTREL

TABLET; ORAL-21

LO/OVRAL

CADENCE HEALTH 0.03MG;0.3MG N017612 001

LOW-OGESTREL-21

MAYNE PHARMA 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

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 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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

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

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

MYLAN

200MG A075800 001 Jan 24, 2003

400MG A075800 002 Jan 24, 2003

ETODOLAC

CAPSULE; ORAL

ETODOLAC

ANI PHARMS INC 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

CHARTWELL MOLECULES 200MG A074842 001 Jul 17, 1997

300MG A074842 002 Jul 17, 1997

ECI PHARMS LLC 300MG A074929 001 Jan 30, 1998

MYLAN 200MG A074932 001 May 16, 1997

200MG A075071 001 Sep 30, 1998

300MG A074932 002 May 16, 1997

300MG A075071 002 Sep 30, 1998

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

CHARTWELL MOLECULES 400MG A074841 001 Jun 27, 1997

ECI PHARMS LLC 400MG A074927 001 Oct 30, 1997

INVATECH 400MG A074839 001 Jul 11, 1997

400MG A074846 001 Feb 28, 1997

IVAX SUB TEVA PHARMS 400MG A074883 001 Feb 28, 1997

500MG A074883 002 Nov 20, 1998

MYLAN 400MG A075012 001 Sep 30, 1998

400MG A075104 001 Feb 06, 1998

500MG A075012 002 Sep 30, 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

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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

6-173(of 430)

** See List Footnote

ETODOLAC

TABLET; ORAL

ETODOLAC

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 INC 400MG

A075943 001 Jul 26, 2002

500MG

A075943 002 Jul 26, 2002

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

LUITPOLD 2MG/ML

A078867 001 Dec 22, 2009

MYLAN LABS LTD 2MG/ML

A078289 001 Jan 02, 2009

PAR STERILE PRODUCTS 2MG/ML

A091297 001 Jun 20, 2012

ETONOGESTREL

IMPLANT; IMPLANTATION

IMPLANON

ORGANON USA INC 68MG/IMPLANT

N021529 001 Jul 17, 2006

ETOPOSIDE

CAPSULE; ORAL

VEPESID

+ DAVA PHARMS INC 50MG

N019557 001 Dec 30, 1986

+ 100MG

N019557 002 Dec 30, 1986

INJECTABLE; INJECTION

ETOPOSIDE

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

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

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** See List Footnote

EVEROLIMUS

TABLET, FOR SUSPENSION;ORAL

EVEROLIMUS

MYLAN	2MG	A210130 001	Apr 19, 2019
	3MG	A210130 002	Apr 19, 2019
	5MG	A210130 003	Apr 19, 2019

EXEMESTANE

TABLET;ORAL

EXEMESTANE

AMNEAL PHARMS	25MG	A206421 001	Dec 28, 2018
MAYNE PHARMA INC	25MG	A208764 001	Aug 08, 2019

EZETIMIBE

TABLET;ORAL

EZETIMIBE

MYLAN	10MG	A201790 001	Apr 26, 2019
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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

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
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INJECTABLE;INJECTION

FAMOTIDINE

APOTEX INC	10MG/ML	A075942 001	Aug 02, 2002
APOTHECON	10MG/ML	A075707 001	Apr 16, 2001
HOSPIRA	10MG/ML	A075705 001	Apr 16, 2001
	10MG/ML	A075870 001	Nov 23, 2001
	10MG/ML	A075905 001	Nov 23, 2001
WEST-WARD PHARMS INT	10MG/ML	A075799 001	Apr 30, 2002

FAMOTIDINE PRESERVATIVE FREE

APOTEX INC	10MG/ML	A076324 001	Nov 27, 2002
APOTHECON	10MG/ML	A075708 001	Apr 16, 2001
HOSPIRA	10MG/ML	A075669 001	Apr 16, 2001
WEST-WARD PHARMS INT	10MG/ML	A075789 001	Apr 30, 2002

FAMOTIDINE PRESERVATIVE FREE (PHARMACY BULK)

APOTEX INC	10MG/ML	A076322 001	Nov 27, 2002
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FAMOTIDINE PRESERVATIVE FREE IN PLASTIC CONTAINER

ABBVIE	0.4MG/ML	A075729 001	Dec 17, 2001
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PEPCID

+	MERCK	10MG/ML **	N019510 001	Nov 04, 1986
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PEPCID PRESERVATIVE FREE

+	MERCK	10MG/ML **	N019510 004	Nov 04, 1986
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PEPCID PRESERVATIVE FREE IN PLASTIC CONTAINER

+	MERCK SHARP DOHME	0.4MG/ML **	N020249 001	Feb 18, 1994
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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	10MG	A075674 001	Dec 21, 2001
	20MG	A075704 001	Apr 16, 2001
	40MG	A075704 002	Apr 16, 2001
MYLAN PHARMS INC	20MG	A075457 001	Apr 18, 2001
	40MG	A075457 002	Apr 18, 2001
PLD ACQUISITIONS	20MG	A075302 001	Apr 16, 2001
	40MG	A075302 002	Apr 16, 2001

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

FAMOTIDINE

TABLET;ORAL

FAMOTIDINE

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 INDUSTRIES	20MG	A075639 002	Dec 12, 2001
	40MG	A075639 001	Dec 12, 2001
WATSON LABS	10MG	A075404 001	Nov 28, 2001
	20MG	A075062 002	Apr 16, 2001
	40MG	A075062 001	Apr 16, 2001

PEPCID

+ VALEANT PHARMS NORTH	20MG	N019462 001	Oct 15, 1986
+	40MG	N019462 002	Oct 15, 1986

TABLET, CHEWABLE;ORAL

PEPCID AC

+ J AND J CONSUMER INC	10MG **	N020801 001	Sep 24, 1998
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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

FELODIPINE

TABLET, EXTENDED RELEASE;ORAL

FELODIPINE

MYLAN	2.5MG	A078855 001	Apr 17, 2008
	5MG	A078855 002	Apr 17, 2008
	10MG	A078855 003	Apr 17, 2008
WOCKHARDT LTD	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 ATLANTIS	87MG	N021695 002	Nov 30, 2004
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FENOFIBRATE (MICRONIZED)

NOVAST LABS	67MG	A207564 001	Apr 19, 2019
	134MG	A207564 002	Apr 19, 2019
	200MG	A207564 003	Apr 19, 2019

LIPIDIL

ABBVIE	100MG	N019304 001	Dec 31, 1993
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LIPOFEN

CIPHER PHARMS INC	100MG	N021612 002	Jan 11, 2006
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TRICOR (MICRONIZED)

+ ABBVIE	67MG **	N019304 002	Feb 09, 1998
+	134MG **	N019304 003	Jun 30, 1999
+	200MG **	N019304 004	Jun 30, 1999

TABLET;ORAL

FENOFIBRATE

MYLAN	107MG	A076520 002	Dec 29, 2005
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TRICOR

+ ABBVIE INC	54MG **	N021203 001	Sep 04, 2001
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+	160MG **	N021203 003	Sep 04, 2001
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TRIGLIDE

SKYE PHARMA AG	50MG	N021350 001	May 07, 2005
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DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

FENOLDOPAM MESYLATE

INJECTABLE; INJECTION

FENOLDOPAM MESYLATE

LUITPOLD	EQ 10MG BASE/ML	A076656	001	Dec 01, 2003
TEVA PARENTERAL	EQ 10MG BASE/ML	A077826	001	Mar 07, 2007

FENOPROFEN CALCIUM

CAPSULE; ORAL

FENOPROFEN CALCIUM

AM THERAP	EQ 200MG BASE	A072307	001	Aug 22, 1988
	EQ 300MG BASE	A072308	001	Aug 22, 1988
AUROLIFE PHARMA LLC	EQ 200MG BASE	A072394	001	Oct 17, 1988
	EQ 300MG BASE	A072395	001	Oct 17, 1988
HALSEY	EQ 200MG BASE	A072355	001	Aug 17, 1988
	EQ 300MG BASE	A072356	001	Aug 17, 1988
PAR PHARM	EQ 200MG BASE	A072437	001	Aug 22, 1988
	EQ 300MG BASE	A072438	001	Aug 22, 1988
QUANTUM PHARMICS	EQ 200MG BASE	A072214	001	Aug 17, 1988
	EQ 300MG BASE	A071738	001	Aug 17, 1988
WARNER CHILCOTT	EQ 200MG BASE	A072946	001	Apr 30, 1991
	EQ 300MG BASE	A072472	001	Apr 30, 1991
WATSON LABS	EQ 200MG BASE	A072294	001	Aug 17, 1988
	EQ 200MG BASE	A072981	001	Aug 19, 1991
	EQ 300MG BASE	A072293	001	Aug 17, 1988
	EQ 300MG BASE	A072982	001	Aug 19, 1991

NALFON

XSPIRE PHARMA	EQ 300MG BASE	N017604	002	
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TABLET; ORAL

FENOPROFEN CALCIUM

AM THERAP	EQ 600MG BASE	A072309	001	Aug 17, 1988
ANI PHARMS INC	EQ 600MG BASE	A072274	001	May 02, 1988
AUROLIFE PHARMA LLC	EQ 600MG BASE	A072396	001	Oct 17, 1988
DAVA PHARMS INC	EQ 600MG BASE	A072326	001	Aug 17, 1988
HALSEY	EQ 600MG BASE	A072357	001	Aug 17, 1988
IVAX SUB TEVA PHARMS	EQ 600MG BASE	A072557	001	Aug 29, 1988
PAR PHARM	EQ 600MG BASE	A072429	001	Aug 17, 1988
QUANTUM PHARMICS	EQ 600MG BASE	A072194	001	Aug 17, 1988
SUN PHARM INDUSTRIES	EQ 600MG BASE	A072902	001	Dec 21, 1990
USL PHARMA	EQ 600MG BASE	A072362	001	Aug 17, 1988
WATSON LABS	EQ 600MG BASE	A072165	001	Aug 17, 1988
	EQ 600MG BASE	A072602	001	Oct 11, 1988
WATSON LABS TEVA	EQ 600MG BASE	A072407	001	Aug 17, 1988

NALFON

DISTA	EQ 600MG BASE	N017710	001	
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FENTANYL

FILM, EXTENDED RELEASE; TRANSDERMAL

FENTANYL-100

ACTAVIS LABS UT INC	100MCG/HR	A076709	004	Aug 20, 2007
MAYNE PHARMA	100MCG/HR	A077062	004	Aug 20, 2007
NOVEN	100MCG/HR	A077775	004	Oct 16, 2009

FENTANYL-25

ACTAVIS LABS UT INC	25MCG/HR	A076709	001	Aug 20, 2007
MAYNE PHARMA	25MCG/HR	A077062	001	Aug 20, 2007
NOVEN	25MCG/HR	A077775	001	Oct 16, 2009

FENTANYL-50

ACTAVIS LABS UT INC	50MCG/HR	A076709	002	Aug 20, 2007
MAYNE PHARMA	50MCG/HR	A077062	002	Aug 20, 2007
NOVEN	50MCG/HR	A077775	002	Oct 16, 2009

FENTANYL-75

ACTAVIS LABS UT INC	75MCG/HR	A076709	003	Aug 20, 2007
MAYNE PHARMA	75MCG/HR	A077062	003	Aug 20, 2007
NOVEN	75MCG/HR	A077775	003	Oct 16, 2009

FENTANYL CITRATE

FILM; BUCCAL

ONSOLIS

BDSI	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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

FENTANYL CITRATEFILM;BUCCAL
ONSOLIS

EQ 1.2MG BASE

N022266 005 Jul 16, 2009

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

SPRAY, METERED;NASAL

LAZANDA

+ BTCP PHARMA

EQ 0.3MG BASE

N022569 003 Dec 21, 2015

TABLET;BUCCAL, SUBLINGUAL

FENTANYL CITRATE

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 0.3MG BASE **

N021947 006 Mar 02, 2007

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

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

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 PYROPHOSPHATE CITRATE

SOLUTION;INTRAVENOUS

TRIFERIC

+ ROCKWELL MEDICAL INC

272MG IRON/50ML (5.44MG IRON/ML)

N206317 002 Sep 04, 2015

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

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

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

ALKEM LABS LTD

4MG

A204827 001 Dec 10, 2015

8MG

A204827 002 Dec 10, 2015

DR REDDYS LABS LTD

4MG

A204975 001 Aug 13, 2019

8MG

A204975 002 Aug 13, 2019

FEXOFENADINE HYDROCHLORIDE

CAPSULE;ORAL

ALLEGRA

SANOFI AVENTIS US

60MG **

N020625 001 Jul 25, 1996

FEXOFENADINE HYDROCHLORIDE

BARR

60MG

A076169 001 Jul 13, 2005

SUSPENSION;ORAL

ALLEGRA

+ SANOFI AVENTIS US

30MG/5ML

N021963 001 Oct 16, 2006

CHILDREN'S ALLEGRA HIVES

+ SANOFI AVENTIS US

30MG/5ML

N201373 002 Jan 24, 2011

CHILDREN'S FEXOFENADINE HYDROCHLORIDE ALLERGY

TARO

30MG/5ML

A208123 001 Nov 09, 2017

CHILDREN'S FEXOFENADINE HYDROCHLORIDE HIVES

TARO

30MG/5ML

A208123 002 Nov 09, 2017

FEXOFENADINE HYDROCHLORIDE

P AND L

30MG/5ML

A201311 001 Jul 25, 2012

TABLET;ORAL

ALLEGRA HIVES

+ SANOFI AVENTIS US

60MG

N020872 008 Jan 24, 2011

+

180MG

N020872 009 Jan 24, 2011

CHILDREN'S ALLEGRA ALLERGY

+ SANOFI AVENTIS US

30MG

N020872 005 Jan 24, 2011

CHILDREN'S ALLEGRA HIVES

+ SANOFI AVENTIS US

30MG

N020872 006 Jan 24, 2011

CHILDREN'S FEXOFENADINE HYDROCHLORIDE ALLERGY

MYLAN

30MG

A077081 004 Jul 21, 2011

CHILDREN'S FEXOFENADINE HYDROCHLORIDE HIVES

MYLAN

30MG

A077081 005 Jul 21, 2011

FEXOFENADINE HYDROCHLORIDE

MYLAN

30MG

A077081 002 Apr 11, 2008

TABLET, ORALLY DISINTEGRATING;ORAL

CHILDREN'S ALLEGRA ALLERGY

+ SANOFI AVENTIS US

30MG

N021909 002 Jan 24, 2011

CHILDREN'S ALLEGRA HIVES

+ SANOFI AVENTIS US

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

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

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

FIBRINOGEN, I-125

INJECTABLE;INJECTION

IBRIN

GE HEALTHCARE 154uCi/VIAL

N017879 001

RADIONUCLIDE-LABELED (125 I) FIBRINOGEN (HUMAN) SENSOR

ABBOTT 140uCi/ML

N017787 001

FINAFLOXACIN

SUSPENSION/DROPS;OTIC

XTORO

+ MERLION PHARMS GMBH 0.3%

N206307 001 Dec 17, 2014

FINASTERIDE

TABLET;ORAL

FINASTERIDE

GEDEON RICHTER USA 5MG

A077251 001 Dec 22, 2006

IVAX SUB TEVA PHARMS 5MG

A076340 001 Jun 19, 2006

MYLAN 1MG

A078161 001 Nov 05, 2013

5MG

A077578 001 Dec 18, 2006

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 INC 50MG

A076030 001 Oct 28, 2002

100MG

A076030 002 Oct 28, 2002

150MG

A076030 003 Oct 28, 2002

APOTEX INC 50MG

A079164 001 Jul 09, 2009

100MG

A079164 002 Jul 09, 2009

150MG

A079164 003 Jul 09, 2009

TAMBOCOR

CNTY LINE PHARMS 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

FLOXURIDINE

INJECTABLE;INJECTION

FLOXURIDINE

AM REGENT 500MG/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

+ 400MG/200ML (2MG/ML)

N019950 005 Jul 08, 1994

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

FLUCONAZOLE

INJECTABLE; INJECTION

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					
MYLAN LABS LTD	200MG/100ML (2MG/ML)	A076888	001	Mar 25, 2005	
MYLAN LABS LTD	400MG/200ML (2MG/ML)	A076888	002	Mar 25, 2005	
FLUCONAZOLE IN SODIUM CHLORIDE 0.9%					
DR REDDYS	200MG/100ML (2MG/ML)	A076653	001	Jul 29, 2004	
DR REDDYS	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	
DR REDDYS	400MG/200ML (2MG/ML)	A076837	002	Jan 13, 2005	
HOSPIRA	200MG/100ML (2MG/ML)	A076617	001	Jul 29, 2004	
HOSPIRA	400MG/200ML (2MG/ML)	A076617	002	Jul 29, 2004	
MYLAN LABS LTD	200MG/100ML (2MG/ML)	A076889	001	Mar 25, 2005	
MYLAN LABS LTD	400MG/200ML (2MG/ML)	A076889	002	Mar 25, 2005	

TABLET; ORAL

FLUCONAZOLE

ANI PHARMS INC	50MG	A076086	001	Jul 29, 2004	
ANI PHARMS INC	100MG	A076086	002	Jul 29, 2004	
ANI PHARMS INC	150MG	A076086	003	Jul 29, 2004	
ANI PHARMS INC	200MG	A076086	004	Jul 29, 2004	
GEDEON RICHTER USA	50MG	A076432	001	Jul 29, 2004	
GEDEON RICHTER USA	100MG	A076432	002	Jul 29, 2004	
GEDEON RICHTER USA	150MG	A076432	003	Jul 29, 2004	
GEDEON RICHTER USA	200MG	A076432	004	Jul 29, 2004	
MYLAN	50MG	A076351	001	Jul 29, 2004	
MYLAN	100MG	A076351	002	Jul 29, 2004	
MYLAN	150MG	A076351	003	Jul 29, 2004	
MYLAN	200MG	A076351	004	Jul 29, 2004	
MYLAN PHARMS INC	50MG	A076042	001	Jul 29, 2004	
MYLAN PHARMS INC	100MG	A076042	002	Jul 29, 2004	
MYLAN PHARMS INC	150MG	A076042	003	Jul 29, 2004	
MYLAN PHARMS INC	200MG	A076042	004	Jul 29, 2004	
PLIVA	50MG	A076424	001	Jul 29, 2004	
PLIVA	100MG	A076424	002	Jul 29, 2004	
PLIVA	150MG	A076424	003	Jul 29, 2004	
PLIVA	200MG	A076424	004	Jul 29, 2004	
RANBAXY LABS LTD	50MG	A076386	001	Jul 29, 2004	
RANBAXY LABS LTD	100MG	A076386	002	Jul 29, 2004	
RANBAXY LABS LTD	150MG	A076386	003	Jul 29, 2004	
RANBAXY LABS LTD	200MG	A076386	004	Jul 29, 2004	
ROXANE	50MG	A076213	001	Jul 29, 2004	
ROXANE	100MG	A076213	002	Jul 29, 2004	
ROXANE	150MG	A076213	003	Jul 29, 2004	
ROXANE	200MG	A076213	004	Jul 29, 2004	

FLUDARABINE PHOSPHATE

INJECTABLE; INJECTION

FLUDARA					
+ GENZYME CORP	50MG/VIAL **	N020038	001	Apr 18, 1991	
FLUDARABINE PHOSPHATE					
MYLAN LABS LTD	50MG/2ML (25MG/ML)	A200647	001	Dec 21, 2011	

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	
+ DOWNSTATE CLINCL	4-90mCi/ML **	N020306	002	Sep 25, 2001	

INJECTABLE; INTRAVENOUS

FLUDEOXYGLUCOSE F18					
+ FEINSTEIN	20-200mCi/ML	N021870	001	Aug 19, 2005	

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

FLUDEOXYGLUCOSE F-18

INJECTABLE; INTRAVENOUS

FLUDEOXYGLUCOSE F18

MIDWEST MEDCL 20-200mCi/ML
WEILL MEDCL COLL 10-100mCi/ML **A203736 001 Nov 19, 2015
N021768 001 Aug 05, 2004FLUDROCORTISONE ACETATE

TABLET; ORAL

FLORINEF

+ CASPER PHARMA LLC 0.1MG **

N010060 001

FLUMAZENIL

INJECTABLE; INJECTION

FLUMAZENIL

BAXTER HLTHCARE CORP 0.5MG/5ML (0.1MG/ML)
1MG/10ML (0.1MG/ML)A076755 002 Oct 12, 2004
A076755 001 Oct 12, 2004DR REDDYS 0.5MG/5ML (0.1MG/ML)
1MG/10ML (0.1MG/ML)A076589 002 Oct 12, 2004
A076589 001 Oct 12, 2004

MYLAN LABS LTD 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 HFA

+ MYLAN SPECIALITY LP 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

FLUCET

ALPHARMA US PHARMS 0.025%

A088360 001 Jan 16, 1984

FLUOCINOLONE ACETONIDE

ACP NIMBLE 0.025%

A089525 001 Jul 26, 1988

ALPHARMA US PHARMS 0.01%

A088361 001 Jan 16, 1984

INVATECH 0.01%

A088047 001 Dec 16, 1982

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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

FLUOCINOLONE ACETONIDE

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

FLUONID

ALLERGAN HERBERT	0.025%	A087157	001	Sep 06, 1984
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FLUOTREX

SAVAGE LABS	0.025%	A088172	001	Mar 09, 1983
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SOLUTION; TOPICAL

FLUOCINOLONE ACETONIDE

ACP NIMBLE	0.01%	A207441	001	Sep 28, 2016
ACTAVIS LABS UT INC	0.01%	A208386	001	Oct 21, 2016
ALPHARMA US PHARMS	0.01%	A087159	001	Jun 16, 1982
BAUSCH AND LOMB	0.01%	A040059	001	Dec 20, 1993
INVATECH	0.01%	A088048	001	Dec 16, 1982
MORTON GROVE	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
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FLUOTREX

SAVAGE LABS	0.01%	A088171	001	Mar 09, 1983
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FLUOCINONIDE

CREAM; TOPICAL

FLUOCINONIDE

AMNEAL PHARMS LLC	0.05%	A210554	001	Aug 21, 2018
PERRIGO NEW YORK	0.05%	A071790	001	Jul 13, 1988

LIDEX

+ CNTY LINE PHARMS	0.05%	N016908	002	
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SOLUTION; TOPICAL

FLUOCINONIDE

TARO	0.05%	A072857	001	Aug 02, 1989
TEVA PHARMS	0.05%	A072522	001	Sep 28, 1990

FLUORESCEIN SODIUM

INJECTABLE; INJECTION

FUNDUSCEIN-25

+ NOVARTIS	25% **	N017869	001	
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FLUOROMETHOLONE

CREAM; TOPICAL

OXYLONE

PHARMACIA AND UPJOHN	0.025%	N011748	001	
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SUSPENSION/DROPS; OPHTHALMIC

FLUOR-OP

NOVARTIS	0.1%	A070185	001	Feb 27, 1986
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FLUOROMETHOLONE ACETATE; TOBRAMYCIN

SUSPENSION/DROPS; OPHTHALMIC

TOBRASONE

EYEVANCE	0.1%; 0.3%	N050628	001	Jul 21, 1989
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FLUOROMETHOLONE; SULFACETAMIDE SODIUM

SUSPENSION/DROPS; OPHTHALMIC

FML-S

ALLERGAN	0.1%; 10%	N019525	001	Sep 29, 1989
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FLUOROURACIL

INJECTABLE; INJECTION

ADRUCIL

PHARMACIA AND UPJOHN	50MG/ML	A081222	001	Jun 28, 1991
+ TEVA PARENTERAL	50MG/ML	N017959	001	
	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
BEDFORD	50MG/ML	A089508	001	Jan 26, 1988

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

FLUOROURACIL

INJECTABLE; INJECTION

FLUOROURACIL

EBEWE PHARMA	500MG/10ML (50MG/ML)	A040772	001	Aug 11, 2008
FRESENIUS KABI USA	50MG/ML	A040291	001	Mar 24, 1999
	50MG/ML	A040379	001	Nov 15, 2000
MARCHAR	50MG/ML	A087791	001	Jan 18, 1983
MYLAN LABS LTD	500MG/10ML (50MG/ML)	A202668	001	Jul 17, 2012
	1GM/20ML (50MG/ML)	A202668	002	Jul 17, 2012
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

SOLUTION; TOPICAL

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

ANI PHARMS INC	EQ 10MG BASE	A076287	001	May 20, 2008
	EQ 20MG BASE	A076287	002	May 20, 2008
BARR	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
CELLTRION	EQ 10MG BASE	A078143	001	Jan 16, 2008
	EQ 20MG BASE	A078143	002	Jan 16, 2008
	EQ 40MG BASE	A078143	003	Jan 16, 2008
CR DOUBLE CRANE	EQ 10MG BASE	A076165	001	Feb 01, 2002
	EQ 20MG BASE	A076165	002	Feb 01, 2002
HERITAGE PHARMS INC	EQ 10MG BASE	A201336	001	Oct 01, 2012
	EQ 20MG BASE	A201336	002	Oct 01, 2012
	EQ 40MG BASE	A201336	003	Oct 01, 2012
MYLAN	EQ 10MG BASE	A075207	001	Jan 30, 2002
	EQ 10MG BASE	A078045	001	Nov 17, 2008
	EQ 20MG BASE	A075207	002	Jan 30, 2002
	EQ 20MG BASE	A078045	002	Nov 17, 2008
	EQ 40MG BASE	A075207	003	May 25, 2007
MYLAN PHARMS INC	EQ 10MG BASE	A075577	001	Jan 29, 2002
	EQ 20MG BASE	A075577	002	Jan 29, 2002
PAR PHARM	EQ 10MG BASE	A076922	001	Dec 16, 2004
	EQ 20MG BASE	A076922	002	Dec 16, 2004
	EQ 40MG BASE	A076922	003	Dec 16, 2004
SANDOZ	EQ 10MG BASE	A077469	001	Nov 17, 2008
	EQ 20MG BASE	A077469	002	Nov 17, 2008

PROZAC

ELI LILLY AND CO	EQ 60MG BASE	N018936	004	Jun 15, 1999
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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
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PROZAC WEEKLY

+	LILLY	EQ 90MG BASE	N021235	001	Feb 26, 2001
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SOLUTION; ORAL

FLUOXETINE HYDROCHLORIDE

ACTAVIS MID ATLANTIC	EQ 20MG BASE/5ML	A075690	001	Jan 31, 2002
AUROBINDO PHARMA LTD	EQ 20MG BASE/5ML	A079209	001	Mar 20, 2009

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

FLUOXETINE HYDROCHLORIDE

SOLUTION; ORAL

FLUOXETINE HYDROCHLORIDE

HI TECH PHARMA	EQ 20MG BASE/5ML	A075525	001	Jun 27, 2002
LANNETT CO INC	EQ 20MG BASE/5ML	A076458	001	May 14, 2004
NOSTRUM LABS INC	EQ 20MG BASE/5ML	A075292	001	Feb 07, 2002

PROZAC

+ LILLY	EQ 20MG BASE/5ML **	N020101	001	Apr 24, 1991
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TABLET; ORAL

FLUOXETINE HYDROCHLORIDE

BARR	EQ 10MG BASE	A075810	001	Feb 01, 2002
FOSUN PHARMA	EQ 10MG BASE	A076024	001	Jan 29, 2002
IVAX SUB TEVA PHARMS	EQ 10MG BASE	A075865	001	Feb 28, 2002
	EQ 40MG BASE	A075865	003	Aug 30, 2004

PROZAC

+ LILLY	EQ 10MG BASE **	N020974	001	Mar 09, 1999
+	EQ 20MG BASE **	N020974	002	Mar 09, 1999

FLUOXYMESTERONE

TABLET; ORAL

ANDROID-F

VALEANT PHARM INTL	10MG	A087196	001	
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FLUOXYMESTERONE

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	
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FLUPHENAZINE ENANTHATE

INJECTABLE; INJECTION

PROLIXIN ENANTHATE

APOTHECON	25MG/ML **	N016110	001	
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FLUPHENAZINE HYDROCHLORIDE

CONCENTRATE; ORAL

FLUPHENAZINE HYDROCHLORIDE

ANI PHARMS INC	5MG/ML	A073058	001	Aug 30, 1991
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PERMITIL

SCHERING	5MG/ML **	N016008	001	
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PROLIXIN

APOTHECON	5MG/ML	A070533	001	Nov 07, 1985
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ELIXIR; ORAL

FLUPHENAZINE HYDROCHLORIDE

ANI PHARMS INC	2.5MG/5ML	A081310	001	Apr 29, 1993
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PROLIXIN

+ APOTHECON	2.5MG/5ML **	N012145	003	
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INJECTABLE; INJECTION

PROLIXIN

APOTHECON	2.5MG/ML **	N011751	005	
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TABLET; ORAL

FLUPHENAZINE HYDROCHLORIDE

WATSON LABS	1MG	A088555	001	Dec 18, 1987
	2.5MG	A088544	001	Dec 18, 1987
	5MG	A088527	001	Dec 18, 1987
	10MG	A088550	001	Dec 18, 1987

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

FLUPHENAZINE HYDROCHLORIDE

TABLET; ORAL

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
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FLUPREDNISOLONE

TABLET; ORAL

ALPHADROL

PHARMACIA AND UPJOHN	1.5MG	N012259	002
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FLURANDRENOLIDE

LOTION; TOPICAL

FLURANDRENOLIDE

ALPHARMA US PHARMS	0.05%	A087203	001	Apr 29, 1982
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OINTMENT; TOPICAL

CORDRAN

+ ALMIRALL	0.025% **	N012806	004
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FLURANDRENOLIDE; NEOMYCIN SULFATE

CREAM; TOPICAL

CORDRAN N

LILLY	0.05%;EQ 3.5MG BASE/GM	N050346	001
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OINTMENT; TOPICAL

CORDRAN N

LILLY	0.05%;EQ 3.5MG BASE/GM	N050345	001
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FLURAZEPAM HYDROCHLORIDE

CAPSULE; ORAL

DALMANE

VALEANT PHARM INTL	15MG **	N016721	001
+	30MG **	N016721	002

FLURAZEPAM HYDROCHLORIDE

AUROLIFE PHARMA LLC	15MG	A071717	002	Jul 31, 1991
	30MG	A071717	001	Jul 31, 1991
HALSEY	15MG	A071808	001	Jan 07, 1988
	30MG	A071809	001	Jan 07, 1988
HERITAGE PHARMA	15MG	A071205	001	Nov 25, 1986
	15MG	A072368	001	Mar 30, 1989
	30MG	A071068	001	Nov 25, 1986
	30MG	A072369	001	Mar 30, 1989
HIKMA INTL PHARMS	15MG	A071107	001	Dec 08, 1986
HIKMA PHARMS	30MG	A071108	001	Dec 08, 1986
PAR PHARM	15MG	A070444	001	Mar 20, 1986
	30MG	A070445	001	Mar 20, 1986
PUREPAC PHARM	15MG	A071927	001	Sep 09, 1987
	30MG	A071551	001	Sep 09, 1987
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
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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

FLURBIPROFEN

TABLET; ORAL

ANSAID

PHARMACIA AND UPJOHN	50MG	N018766 002	Oct 31, 1988
	100MG	N018766 003	Oct 31, 1988

FLURBIPROFEN

AUROLIFE PHARMA LLC	50MG	A074448 001	Jul 28, 1995
	100MG	A074448 002	Jul 28, 1995
IVAX SUB TEVA PHARMS	50MG	A074411 001	May 31, 1995
	100MG	A074411 002	May 31, 1995
PLIVA	50MG	A074647 001	Apr 01, 1997
	100MG	A074647 002	Apr 01, 1997
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

FLUTAMIDE

CAPSULE; ORAL

EULEXIN

+ SCHERING	125MG	N018554 001	Jan 27, 1989
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FLUTAMIDE

ACTAVIS LABS FL INC	125MG	A075820 001	Sep 18, 2001
MYLAN	125MG	A076224 001	May 09, 2003
YAOPHARMA CO LTD	125MG	A075818 001	Sep 18, 2001

FLUTEMETAMOL F-18

INJECTABLE; INTRAVENOUS

VIZAMYL

+ GE HEALTHCARE	40.5mCi/10ML (4.05mCi/ML)	N203137 001	Oct 25, 2013
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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
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FLUTICASONE PROPIONATE

NESHER PHARMS	0.05%	A076865 001	Sep 10, 2004
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OINTMENT; TOPICAL

CUTIVATE

+ FOUGERA PHARMS	0.005%	N019957 001	Dec 14, 1990
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FLUTICASONE PROPIONATE

FOUGERA PHARMS	0.005%	A076300 001	May 14, 2004
TARO PHARM INDS	0.005%	A077145 001	Jun 14, 2005

POWDER; INHALATION

ARMONAIR RESPICLICK

+ TEVA PHARM	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

+ GLAXOSMITHKLINE	0.05MG/SPRAY **	N020121 001	Oct 19, 1994
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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

MYLAN	EQ 80MG BASE	A202458 001	Sep 11, 2015
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Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

FLUVOXAMINE MALEATE

CAPSULE, EXTENDED RELEASE;ORAL

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 INC	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	
ECI PHARMS LLC	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	25MG	A075889 001	Nov 29, 2000	
	50MG	A075889 002	Nov 29, 2000	
	50MG	A075950 001	Oct 15, 2001	
	100MG	A075889 003	Nov 29, 2000	
	100MG	A075950 002	Oct 15, 2001	
SUN PHARM INDUSTRIES	25MG	A076125 001	Apr 29, 2002	
	50MG	A076125 002	Apr 29, 2002	
	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	
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
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FOLVITE

+	WYETH PHARMS INC	5MG/ML	N005897 008
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TABLET;ORAL

FOLIC ACID

BARR	1MG	A089177 001	Jan 08, 1986
CONTRACT PHARMACAL	1MG	A085061 001	
EVERYLIFE	1MG	A080755 001	
HALSEY	1MG	A083598 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	

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

FOLIC ACID

TABLET; ORAL

FOLIC ACID

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	

FOLLITROPIN ALFA/BETA

INJECTABLE; INTRAMUSCULAR, SUBCUTANEOUS

FOLLISTIM

ORGANON USA INC	75 IU/VIAL	N020582 001	Sep 29, 1997
	150 IU/VIAL	N020582 002	Sep 29, 1997

INJECTABLE; SUBCUTANEOUS

FOLLISTIM AQ

ORGANON USA INC	75 IU/0.5ML	N021273 001	Aug 26, 2005
	150 IU/0.18ML	N021211 003	Feb 11, 2004
	150 IU/0.5ML	N021273 002	Aug 26, 2005

GONAL-F

EMD SERONO	37.5 IU/VIAL	N020378 003	May 25, 2000
	37.5 IU/VIAL	N021765 001	Mar 25, 2004
	75 IU/VIAL	N020378 001	Sep 29, 1997
	150 IU/VIAL	N020378 002	Sep 29, 1997
	150 IU/VIAL	N021765 003	Mar 25, 2004

FOMEPIZOLE

INJECTABLE; INJECTION

ANTIZOL

+ PAR PHARM INC	1.5GM/1.5ML (1GM/ML)	N020696 001	Dec 04, 1997
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FOMEPIZOLE

MYLAN INSTITUTIONAL	1.5GM/1.5ML (1GM/ML)	A079033 001	Apr 07, 2009
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FOMIVIRSEN SODIUM

INJECTABLE; INJECTION

VITRAVENE PRESERVATIVE FREE

NOVARTIS	6.6MG/ML	N020961 001	Aug 26, 1998
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FORMOTEROL FUMARATE

POWDER; INHALATION

FORADIL

+ NOVARTIS	0.012MG/INH	N020831 001	Feb 16, 2001
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FORADIL CERTIHALER

NOVARTIS	0.0085MG/INH	N021592 001	Dec 15, 2006
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FOSAPREPITANT DIMEGLUMINE

POWDER; INTRAVENOUS

EMEND

+ MERCK AND CO INC	EQ 115MG BASE/VIAL **	N022023 001	Jan 25, 2008
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FOSCARNET SODIUM

INJECTABLE; INJECTION

FOSCARNET SODIUM

HOSPIRA	2.4GM/100ML	A077174 001	May 31, 2005
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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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

FOSINOPRIL SODIUM

TABLET; ORAL

FOSINOPRIL SODIUM

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 INC	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
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

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
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

FOSPROPOFOL DISODIUM

SOLUTION; INTRAVENOUS

LUSEDRA

EISAI INC	1050MG/30ML (35MG/ML)	N022244	001	Dec 12, 2008
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FURAZOLIDONE

SUSPENSION; ORAL

FUROXONE

SHIRE	50MG/15ML	N011323	002	
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TABLET; ORAL

FUROXONE

SHIRE	100MG	N011270	002	
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FUROSEMIDE

INJECTABLE; INJECTION

FUROSEMIDE

ABRAXIS PHARM	10MG/ML	N018507	001	Jul 30, 1982
	10MG/ML	N019036	001	Aug 13, 1984
ACCORD HLTHCARE	10MG/ML	A070017	001	Dec 15, 1986
+ AM REGENT	10MG/ML **	N018579	001	Nov 30, 1983
ASTRAZENECA	10MG/ML	A070014	001	Sep 09, 1985
HOSPIRA	10MG/ML	A070578	001	Jul 08, 1987
	10MG/ML	A072080	001	Aug 13, 1991
	10MG/ML	A074337	001	Oct 31, 1994
IGI LABS INC	10MG/ML	A070095	001	Sep 09, 1985
	10MG/ML	A070096	001	Sep 09, 1985
	10MG/ML	N018025	001	
INTL MEDICATION	10MG/ML	A074017	001	Jun 30, 1994
MARSAM PHARMS LLC	10MG/ML	A070023	001	Feb 05, 1986
SMITH AND NEPHEW	10MG/ML	A070078	001	Feb 05, 1986
	10MG/ML	N018420	001	Feb 26, 1982
WARNER CHILCOTT	10MG/ML	A070019	001	Sep 22, 1986
WATSON LABS	10MG/ML	A070604	001	Jan 02, 1987
	10MG/ML	A071439	001	Sep 14, 1990
WEST-WARD PHARMS INT	10MG/ML			

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

FUROSEMIDE

INJECTABLE; INJECTION

FUROSEMIDE

	10MG/ML	N018267 001	
WYETH AYERST	10MG/ML	N018670 001	Jul 20, 1982
LASIX			
+ SANOFI AVENTIS US	10MG/ML **	N016363 001	

SOLUTION; ORAL

LASIX

SANOFI AVENTIS US	10MG/ML	N017688 001	
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TABLET; ORAL

FUROSEMIDE

ANI PHARMS INC	20MG	A071379 001	Jan 02, 1987
	40MG	A070413 001	Feb 26, 1986
	80MG	A071594 001	Feb 09, 1988
AVET	20MG	N018413 001	Nov 30, 1983
	40MG	N018413 002	Nov 30, 1983
DAVA PHARMS INC	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
SANDOZ	40MG	N018750 002	Jul 30, 1984
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

HIKMA	100MG	A078150 001	Sep 25, 2007
	300MG	A078150 002	Sep 25, 2007
	400MG	A078150 003	Sep 25, 2007
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
	300MG	A076606 002	Oct 07, 2005
	400MG	A076606 003	Oct 07, 2005
SUN PHARM INDUSTRIES	100MG	A076537 001	Jun 30, 2005
	300MG	A076537 002	Jun 30, 2005
	400MG	A076537 003	Jun 30, 2005
WATSON LABS	100MG	A075485 003	May 11, 2007
	300MG	A075485 002	May 11, 2007
	400MG	A075485 001	May 11, 2007

TABLET; ORAL

GABAPENTIN

HIKMA PHARMS	600MG	A078782 001	Jul 21, 2011
	800MG	A078782 002	Jul 21, 2011
INVATECH	600MG	A076877 001	Jul 06, 2006
	800MG	A076877 002	Jul 06, 2006

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

6-191(of 430)

** See List Footnote

GABAPENTIN

TABLET; ORAL

GABAPENTIN

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
RANBAXY	600MG	A076605 001	Sep 14, 2005
	800MG	A076605 002	Sep 14, 2005
SANDOZ	600MG	A076120 001	Jan 27, 2006
	800MG	A076120 002	Jan 27, 2006
TEVA	600MG	A075827 001	Dec 15, 2004
	800MG	A075827 002	Dec 15, 2004

GADODIAMIDE

INJECTABLE; INJECTION

OMNISCAN

GE HEALTHCARE	14.35GM/50ML (287MG/ML)	N022066 001	Sep 05, 2007
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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

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
MYLAN	EQ 8MG BASE	A090900 001	Jan 24, 2011
	EQ 16MG BASE	A090900 002	Jan 24, 2011
	EQ 24MG BASE	A090900 003	Jan 24, 2011

SOLUTION; ORAL

RAZADYNE

JANSSEN PHARMS	4MG/ML **	N021224 001	Jun 22, 2001
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TABLET; ORAL

GALANTAMINE HYDROBROMIDE

HERITAGE PHARMA	EQ 4MG BASE	A077585 001	Sep 15, 2009
	EQ 4MG BASE	A077587 001	Jul 09, 2009
	EQ 8MG BASE	A077585 002	Sep 15, 2009
	EQ 8MG BASE	A077587 002	Jul 09, 2009
	EQ 12MG BASE	A077585 003	Sep 15, 2009
	EQ 12MG BASE	A077587 003	Jul 09, 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 4MG BASE	A077603 001	Aug 28, 2008
	EQ 8MG BASE	A077590 002	May 29, 2009
	EQ 8MG BASE	A077603 002	Aug 28, 2008

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

6-192(of 430)

** See List Footnote

GALANTAMINE HYDROBROMIDE

TABLET; ORAL

GALANTAMINE HYDROBROMIDE

EQ 12MG BASE

A077590 003 May 29, 2009

EQ 12MG BASE

A077603 003 Aug 28, 2008

GALLAMINE TRIETHIODIDE

INJECTABLE; INJECTION

FLAXEDIL

DAVIS AND GECK

20MG/ML

N007842 001

100MG/ML

N007842 002

GALLIUM CITRATE GA-67

INJECTABLE; INJECTION

GALLIUM CITRATE GA 67

GE HEALTHCARE

1mCi/ML

N017700 001

NEOSCAN

GE HEALTHCARE

2mCi/ML

N017655 001

GALLIUM NITRATE

INJECTABLE; INJECTION

GANITE

CHEPLAPHARM

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

GANCICLOVIR SODIUM

AM REGENT

EQ 500MG BASE/VIAL

A202624 001 Sep 18, 2013

CUSTOPHARM INC

EQ 500MG BASE/VIAL

A212001 001 Jun 20, 2019

WEST-WARD PHARMS INT

EQ 500MG BASE/VIAL

A076222 001 Jul 16, 2003

GATIFLOXACIN

SOLUTION/DROPS; OPHTHALMIC

GATIFLOXACIN

APOTEX INC

0.3%

A079084 001 Aug 19, 2011

GEFITINIB

TABLET; ORAL

IRESSA

ASTRAZENECA

250MG

N021399 001 May 05, 2003

GEMCITABINE HYDROCHLORIDE

INJECTABLE; INJECTION

GEMCITABINE HYDROCHLORIDE

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

HAMELN RDS GMBH

EQ 200MG BASE/VIAL

A090663 001 Sep 10, 2012

EQ 1GM BASE/VIAL

A090663 002 Sep 10, 2012

INGENUS PHARMS LLC

200MG/5.26ML (38MG/ML)

A210383 001 Feb 14, 2019

1GM/26.3ML (38MG/ML)

A210383 002 Feb 14, 2019

2GM/52.6ML (38MG/ML)

A210383 003 Feb 14, 2019

SAGENT PHARMS

EQ 200MG BASE/VIAL

A091597 001 May 07, 2013

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

6-193(of 430)

** See List Footnote

GEMCITABINE HYDROCHLORIDE

INJECTABLE; INJECTION

GEMCITABINE HYDROCHLORIDE

EQ 1GM BASE/VIAL A091597 002 May 07, 2013

GEMFIBROZIL

CAPSULE; ORAL

GEMFIBROZIL

MYLAN

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

MYLAN

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

YAOPHARMA CO LTD

600MG

A074615 001 Sep 29, 1995

GEMTUZUMAB OZOGAMICIN

INJECTABLE; INJECTION

MYLOTARG

WYETH PHARMS INC

5MG/VIAL

N021174 001 May 17, 2000

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

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

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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

6-194(of 430)

** See List Footnote

GENTAMICIN SULFATE

INJECTABLE; INJECTION

GENTAMICIN SULFATE

	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	
WEST-WARD PHARMS INT	EQ 10MG BASE/ML	A062251 002	
	EQ 40MG BASE/ML	A062251 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
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
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
PHARMADERM	EQ 0.1% BASE	A062534 001	Oct 10, 1984

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

6-195(of 430)

** See List Footnote

GENTAMICIN SULFATE

SOLUTION/DROPS;OPHTHALMIC

GARAMYCIN

+ SCHERING

EQ 0.3% BASE **

N050039 002

GENTACIDIN

NOVARTIS

EQ 0.3% BASE

A062480 001 Mar 30, 1984

GENTAFAIR

PHARMAFAIR

EQ 0.3% BASE

A062440 001 May 03, 1983

GENTAMICIN SULFATE

ALCON PHARMS LTD

EQ 0.3% BASE

A062523 001 Nov 25, 1985

PACO

EQ 3MG BASE/ML

A062932 001 Nov 07, 1988

GENTIAN VIOLET

SUPPOSITORY;VAGINAL

GVS

SAVAGE LABS

0.4%

A083513 001

TAMPON;VAGINAL

GENAPAX

KEY PHARMS

5MG

A085017 001

GLATIRAMER ACETATE

FOR 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

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

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

GLIPIZIDE

TABLET;ORAL

GLIPIZIDE

ANI PHARMS INC	5MG	A074387 001	Mar 04, 1996
	5MG	A074497 001	Aug 31, 1995
	10MG	A074387 002	Mar 04, 1996
	10MG	A074497 002	Aug 31, 1995
BARR LABS INC	5MG	A074619 001	Apr 04, 1997
	10MG	A074619 002	Apr 04, 1997
BEXIMCO PHARMS USA	5MG	A074542 001	Jun 20, 1995
	10MG	A074542 002	Jun 20, 1995
MYLAN	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
WATSON LABS	5MG	A074370 001	Nov 22, 1994
	10MG	A074370 002	Nov 22, 1994

GLUCOTROL

PFIZER	2.5MG	N017783 003	May 11, 1993
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TABLET, EXTENDED RELEASE;ORAL

GLIPIZIDE

MYLAN	2.5MG	A202298 001	May 19, 2015
	5MG	A202298 002	May 19, 2015
	10MG	A202298 003	May 19, 2015

GLIPIZIDE; METFORMIN HYDROCHLORIDE

TABLET;ORAL

GLIPIZIDE AND METFORMIN HYDROCHLORIDE

MYLAN	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 HYDROCHLORIDE

INJECTABLE; INJECTION

GLUCAGON

+	LILLY	EQ 1MG BASE/VIAL **	N012122 001
+		EQ 10MG BASE/VIAL **	N012122 002

GLUTETHIMIDE

CAPSULE;ORAL

DORIDEN

SANOFI AVENTIS US	500MG	N009519 008
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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	

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
ORIENT PHARMA CO LTD	1.25MG	A206483 001	Feb 22, 2019

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

6-197(of 430)

** See List Footnote

GLYBURIDE

TABLET; ORAL

GLYBURIDE

2.5MG

A206483 002 Feb 22, 2019

5MG

A206483 003 Feb 22, 2019

GLYBURIDE (MICRONIZED)

MYLAN

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

YAOPHARMA CO LTD

1.5MG

A075174 001 Jun 22, 1998

3MG

A075174 002 Jun 22, 1998

GLYNASE

+ PHARMACIA AND UPJOHN 4.5MG **

N020051 003 Sep 24, 1993

MICRONASE

+ PHARMACIA AND UPJOHN 1.25MG **

N017498 001 May 01, 1984

2.5MG

N017498 002 May 01, 1984

+ 5MG **

N017498 003 May 01, 1984

GLYBURIDE; METFORMIN HYDROCHLORIDE

TABLET; ORAL

GLUCOVANCE

+ BRISTOL MYERS SQUIBB 1.25MG; 250MG **

N021178 001 Jul 31, 2000

+ 2.5MG; 500MG **

N021178 002 Jul 31, 2000

+ 5MG; 500MG **

N021178 003 Jul 31, 2000

GLYBURIDE AND METFORMIN HYDROCHLORIDE

HERITAGE PHARMS INC 1.25MG; 250MG

A079009 001 Jun 03, 2009

2.5MG; 500MG

A079009 002 Jun 03, 2009

5MG; 500MG

A079009 003 Jun 03, 2009

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

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

HOSPIRA 0.2MG/ML

A089393 001 Jun 15, 1988

TEVA PARENTERAL 0.2MG/ML

A081169 001 Sep 10, 1991

WATSON LABS 0.2MG/ML

A086947 001 Jun 24, 1983

ROBINUL

+ HIKMA 0.2MG/ML **

N017558 001

ROBINS AH 0.2MG/ML

N014764 001

TABLET; ORAL

GLYCOPYRROLATE

HIKMA INTL PHARMS 1MG

A040836 001 Mar 05, 2009

2MG

A040836 002 Mar 05, 2009

RENATA 1MG

A040568 001 Dec 22, 2004

2MG

A040568 002 Dec 22, 2004

WATSON LABS 1MG

A085562 001

1MG

A086902 001

2MG

A085563 001

2MG

A086178 001

2MG

A086900 001

ROBINUL

+ CASPER PHARMA LLC 1MG **

N012827 001

ROBINUL FORTE

+ CASPER PHARMA LLC 2MG **

N012827 002

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

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

WEST-WARD PHARMS INT	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

GONADOTROPIN, CHORIONIC

INJECTABLE; INJECTION

A.P.L.

FERRING	5,000 UNITS/VIAL	N017055 001	
	10,000 UNITS/VIAL	N017055 002	
	20,000 UNITS/VIAL	N017055 003	

CHORIONIC GONADOTROPIN

BEL MAR	5,000 UNITS/VIAL	N017054 001	
	10,000 UNITS/VIAL	N017054 002	
FERRING	2,000 UNITS/VIAL	N017016 009	Dec 27, 1984
	2,000 UNITS/VIAL	N017016 011	Feb 16, 1990
	15,000 UNITS/VIAL	N017016 010	Feb 15, 1985
	20,000 UNITS/VIAL	N017016 004	
FRESENIUS KABI USA	5,000 UNITS/VIAL	N017067 001	
	15,000 UNITS/VIAL	N017067 004	
	20,000 UNITS/VIAL	N017067 003	

FOLLUTEIN

BRISTOL MYERS SQUIBB	10,000 UNITS/VIAL	N017056 001	
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GRAMICIDIN; NEOMYCIN SULFATE; POLYMYXIN B SULFATE

SOLUTION/DROPS; OPHTHALMIC

NEO-POLYCYCIN

DOW PHARM	0.025MG/ML; EQ 1.75MG BASE/ML; 10,000 UNITS/ML	A060427 001	
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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
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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
SANDOZ INC	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
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
+ ROCHE	EQ 1MG BASE/ML (EQ 1MG BASE/ML) **	N020239 004	Mar 11, 1994
+ ROCHE	EQ 3MG BASE/ML **	N020239 001	Dec 29, 1993
+ ROCHE	EQ 4MG BASE/4ML (EQ 1MG BASE/ML) **	N020239 002	Mar 11, 1994

SOLUTION; ORAL

GRANISOL

PEDIATRAX	EQ 2MG BASE/10ML	A078334 001	Feb 28, 2008
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KYTRIL

+ ROCHE	EQ 2MG BASE/10ML **	N021238 001	Jun 27, 2001
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TABLET; ORAL

GRANISETRON HYDROCHLORIDE

BARR	EQ 1MG BASE	A078221 001	Dec 31, 2007
EPIC PHARMA LLC	EQ 1MG BASE	A078260 001	Dec 31, 2007
MYLAN	EQ 1MG BASE	A078725 001	Jan 30, 2008

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

6-199(of 430)

** See List Footnote

GRANISETRON HYDROCHLORIDE

TABLET;ORAL

GRANISETRON HYDROCHLORIDE

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

FULVICIN-U/F

CHARTWELL RX

250MG

A060569 002

500MG

A060569 001

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

GRISEOFULVIN, ULTRAMICROCRYSTALLINE

TABLET;ORAL

FULVICIN P/G

CHARTWELL RX

125MG

A061996 001

250MG

A061996 002

FULVICIN P/G 165

CHARTWELL RX

165MG

A061996 003 Apr 06, 1982

FULVICIN P/G 330

CHARTWELL RX

330MG

A061996 004 Apr 06, 1982

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

TABLET, EXTENDED RELEASE;ORAL

GUAIFENESIN

OHM LABS INC

600MG

A209254 001 Jul 16, 2018

1.2GM

A209254 002 Jul 16, 2018

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

6-200(of 430)

** See List Footnote

GUAIFENESIN; HYDROCODONE BITARTRATE

SOLUTION; ORAL

FLOWTUSS

BKK PHARMS 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

+ BKK PHARMS 200MG/5ML; 2.5MG/5ML; 30MG/5ML N022279 001 May 14, 2015

GUANABENZ ACETATE

TABLET; ORAL

GUANABENZ ACETATE

ANI PHARMS INC 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

WATSON LABS EQ 4MG BASE A074025 001 Feb 28, 1994

EQ 8MG BASE A074025 002 Feb 28, 1994

YAOPHARMA CO LTD EQ 4MG BASE A074517 001 Sep 30, 1998

EQ 8MG BASE A074517 002 Sep 30, 1998

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

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

MYLAN 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

DISCONTINUED DRUG PRODUCT LIST

6-201(of 430)

** See List Footnote

HALAZEPAM

TABLET;ORAL

PAXIPAM

SCHERING	20MG	N017736 003
	40MG	N017736 004

HALCINONIDE

CREAM;TOPICAL

HALOG

WESTWOOD SQUIBB	0.025%	N017818 001
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HALOG-E

SUN PHARM INDS INC	0.1%	N018234 001
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OINTMENT;TOPICAL

HALOG

BRISTOL MYERS SQUIBB	0.025%	N018125 001
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SOLUTION;TOPICAL

HALOG

SUN PHARM INDS INC	0.1%	N017823 001
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HALOBETASOL PROPIONATE

CREAM;TOPICAL

ULTRAVATE

+ SUN PHARM INDS INC	0.05%	N019967 001	Dec 27, 1990
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OINTMENT;TOPICAL

HALOBETASOL PROPIONATE

FOUGERA PHARMS	0.05%	A076903 001	Dec 16, 2004
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HALOFANTRINE HYDROCHLORIDE

TABLET;ORAL

HALFAN

GLAXOSMITHKLINE	250MG	N020250 001	Jul 24, 1992
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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
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HALOPERIDOL

CYCLE PHARMS LTD	0.5MG	A071128 001	Feb 17, 1987
	1MG	A071129 001	Feb 17, 1987
	2MG	A071130 001	Feb 17, 1987
	5MG	A071131 001	Feb 17, 1987
	10MG	A071132 001	May 12, 1987
	20MG	A071133 001	May 12, 1987
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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

HALOPERIDOL

TABLET; ORAL

HALOPERIDOL

	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
SANDOZ	2MG	A071208	001	Nov 17, 1986
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
VINTAGE	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
	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 INC	EQ 50MG BASE/ML	A076463	001	Jun 24, 2005
	EQ 100MG BASE/ML	A076463	002	Jun 24, 2005

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				
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
FOSUN PHARMA	EQ 5MG BASE/ML	A076464	001	Sep 29, 2004
FRESENIUS KABI USA	EQ 5MG BASE/ML	A210356	001	Jul 01, 2019
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
WATSON LABS	EQ 5MG BASE/ML	A070713	001	May 17, 1988

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

HALOPERIDOL LACTATE

INJECTABLE; INJECTION

HALOPERIDOL

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

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

AKORN 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

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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

HEPARIN SODIUM

INJECTABLE; INJECTION

HEPARIN SODIUM

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	
FRESENIUS KABI USA	1,000 UNITS/ML	N017651 005	
	5,000 UNITS/ML	N017029 002	
	10,000 UNITS/ML	N017651 003	
	20,000 UNITS/ML	N017651 008	
HIKMA	5,000 UNITS/0.5ML	N017037 013	Apr 07, 1986
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
ORGANON USA INC	1,000 UNITS/ML	N000552 008	
	5,000 UNITS/ML	N000552 009	
	10,000 UNITS/ML	N000552 010	
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	
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
+ WEST-WARD PHARMS INT	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	
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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

HEPARIN SODIUM

INJECTABLE; INJECTION

HEPARIN SODIUM 10,000 UNITS	IN SODIUM CHLORIDE 0.9%			
	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
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
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				
ORGANON USA INC	100 UNITS/ML		N000552 007	
LIQUAEMIN SODIUM				
ORGANON USA 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	

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

HEPARIN SODIUM

INJECTABLE; INJECTION

LIQUAEMIN SODIUM

	40,000 UNITS/ML	N000552 002	
LIQUAEMIN SODIUM PRESERVATIVE FREE			
ORGANON USA 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	
	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	
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TURGEX

XTTRIUM	3%	N018375 001	
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EMULSION; TOPICAL

HEXA-GERM

HUNTINGTON LABS	3%	N017411 001	
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PHISOHEX

SANOFI AVENTIS US	3%	N006882 001	
	3%	N008402 001	

SOY-DOME

BAYER PHARMS	3%	N017405 001	
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TURGEX

XTTRIUM	3%	N019055 001	Nov 30, 1984
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SOAP; TOPICAL

GAMOPHEN

ARBROOK	2%	N006270 003	
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SOLUTION; TOPICAL

DIAL

DIAL	0.25%	N017421 002	
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GERMA-MEDICA

HUNTINGTON LABS	1%	N017412 001	
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GERMA-MEDICA "MG"

HUNTINGTON LABS	0.25%	N017412 002	
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SEPTI-SOFT

CALGON	0.25%	N017460 001	
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SEPTISOL

VESTAL LABS	0.25%	N017423 001	
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SPONGE; TOPICAL

E-Z SCRUB

BECTON DICKINSON	450MG	N017452 001	
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HEXASCRUB

PROF DSPLS	3%	N018363 001	
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Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

6-207(of 430)

** See List Footnote

HEXACHLOROPHENE

SPONGE; TOPICAL

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

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

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

BRECKENRIDGE 1.5MG/5ML; 5MG/5ML

A210663 001 Jun 11, 2019

IVAX SUB TEVA PHARMS 1.5MG/5ML; 5MG/5ML

A040285 001 Jul 19, 1999

TORRENT 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

HYCODAN

+ GENUS 1.5MG; 5MG **

N005213 001 Jul 26, 1988

TUSSIGON

KING PHARMS 1.5MG; 5MG

A088508 001 Jul 30, 1985

HYALURONIDASE

INJECTABLE; INJECTION

HYDASE

AKORN INC 150 UNITS/ML

N021716 001 Oct 25, 2005

VITRASE

BAUSCH AND LOMB 6,200 UNITS/VIAL

N021640 001 May 05, 2004

WYDASE

BAXTER HLTHCARE 150 UNITS/ML **

N006343 002

150 UNITS/VIAL **

N006343 006

1,500 UNITS/VIAL **

N006343 005

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

6-208(of 430)

** See List Footnote

HYDRALAZINE HYDROCHLORIDE

INJECTABLE; INJECTION

APRESOLINE

+ NOVARTIS 20MG/ML ** N008303 003

HYDRALAZINE HYDROCHLORIDE

ABRAXIS PHARM 20MG/ML A089532 001 Aug 11, 1987

AM REGENT 20MG/ML A040136 001 Jun 30, 1997

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

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

HERITAGE PHARMS INC 10MG A040858 001 Feb 26, 2010

25MG A040858 002 Feb 26, 2010

50MG A040858 003 Feb 26, 2010

100MG A040858 004 Feb 26, 2010

IMPAX LABS 25MG A084922 001

50MG A084923 001

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

MYLAN 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

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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

6-209(of 430)

** See List Footnote

HYDRALAZINE HYDROCHLORIDE

TABLET; ORAL

HYDRALAZINE HYDROCHLORIDE

	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

HYDRALAZINE HYDROCHLORIDE; HYDROCHLOROTHIAZIDE

CAPSULE; ORAL

APRESAZIDE

NOVARTIS	25MG; 25MG	A084735 001	
	50MG; 50MG	A084810 001	
	100MG; 50MG	A084811 001	

HYDRA-ZIDE

PAR PHARM	100MG; 50MG	A088961 001	Oct 21, 1985
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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
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HYDRALAZINE HYDROCHLORIDE W/ HYDROCHLOROTHIAZIDE 25/25

IVAX PHARMS	25MG; 25MG	A088356 001	Apr 10, 1984
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HYDRALAZINE HYDROCHLORIDE W/ HYDROCHLOROTHIAZIDE 50/50

IVAX PHARMS	50MG; 50MG	A088357 001	Apr 10, 1984
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TABLET; ORAL

APRESOLINE-ESIDRIX

NOVARTIS	25MG; 15MG	N012026 002	
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HYDRALAZINE AND HYDROCHLOROTHIAZIDE

WATSON LABS	25MG; 15MG	A085827 001	
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HYDROCHLOROTHIAZIDE W/ HYDRALAZINE

WATSON LABS	25MG; 15MG	A085373 001	
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HYDRALAZINE HYDROCHLORIDE; HYDROCHLOROTHIAZIDE; RESERPINE

TABLET; ORAL

CAM-AP-ES

CHARTWELL RX	25MG; 15MG; 0.1MG	A084897 001	
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HYDRALAZINE HYDROCHLORIDE, HYDROCHLOROTHIAZIDE AND RESERPINE

IVAX SUB TEVA PHARMS	25MG; 15MG; 0.1MG	A084291 001	
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HYDRALAZINE HYDROCHLORIDE-HYDROCHLOROTHIAZIDE-RESERPINE

MYLAN	25MG; 15MG; 0.1MG	A087085 001	
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HYDRALAZINE, HYDROCHLOROTHIAZIDE W/ RESERPINE

WATSON LABS	25MG; 15MG; 0.1MG	A085771 001	
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HYDRAP-ES

SANDOZ	25MG; 15MG; 0.1MG	A084876 001	
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HYDROCHLOROTHIAZIDE W/ RESERPINE AND HYDRALAZINE

WATSON LABS	25MG; 15MG; 0.1MG	A083770 001	
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HYDROSERPINE PLUS (R-H-H)

IVAX SUB TEVA PHARMS	25MG; 15MG; 0.1MG	A083877 001	
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RESERPINE, HYDRALAZINE HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE

SOLVAY	25MG; 15MG; 0.1MG	A088376 001	Oct 28, 1983
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SUN PHARM INDUSTRIES	25MG; 15MG; 0.1MG	A088570 001	Apr 10, 1984
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WATSON LABS	25MG; 15MG; 0.1MG	A085549 001	
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	25MG; 15MG; 0.1MG	A087556 001	
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RESERPINE, HYDROCHLOROTHIAZIDE, AND HYDRALAZINE HYDROCHLORIDE

LEDERLE	25MG; 15MG; 0.1MG	A087709 001	May 13, 1982
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Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

6-210(of 430)

** See List Footnote

HYDRALAZINE HYDROCHLORIDE; HYDROCHLOROTHIAZIDE; RESERPINE

TABLET; ORAL

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

HYDROCHLOROTHIAZIDE

CAPSULE; ORAL

HYDROCHLOROTHIAZIDE

APOTEX 12.5MG A078389 001 May 16, 2008

HIKMA INTL PHARMS 12.5MG A077885 001 Nov 26, 2007

LANNETT CO INC 12.5MG A091662 001 Jan 27, 2012

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

50MG A083965 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

IMPAX LABS 25MG A084029 001

50MG A083607 002

100MG A085098 001

INWOOD LABS 25MG A084776 001

25MG A085067 001

50MG A084776 002

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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

HYDROCHLOROTHIAZIDE

TABLET; ORAL

HYDROCHLOROTHIAZIDE

	50MG	A086192 002	
MYLAN	25MG	A084880 001	
	50MG	A085112 001	
MYLAN PHARMS INC	12.5MG	A040770 001	Jan 23, 2007
	25MG	A040735 002	Jan 23, 2007
	50MG	A040735 003	Jan 23, 2007
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	
YAOPHARMA CO LTD	25MG	A087565 001	Mar 09, 1982
	50MG	A084912 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
MYLAN PHARMS INC	12.5MG; 150MG	A077969 001	Sep 27, 2012

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

HYDROCHLOROTHIAZIDE; IRBESARTAN

TABLET; ORAL

IRBESARTAN AND HYDROCHLOROTHIAZIDE

	12.5MG; 300MG	A077969 002	Sep 27, 2012
	25MG; 300MG	A077969 003	Jul 20, 2016
TEVA	25MG; 300MG	A077369 003	Mar 30, 2012
WATSON LABS INC	12.5MG; 150MG	A091539 001	Oct 22, 2012
	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

COREPHARMA	12.5MG; 10MG	A076674 001	Oct 05, 2004
	12.5MG; 20MG	A076674 002	Oct 05, 2004
	25MG; 20MG	A076674 003	Oct 05, 2004
HERITAGE PHARMA	12.5MG; 10MG	A075776 001	Jul 01, 2002
	12.5MG; 20MG	A075776 002	Jul 01, 2002
	25MG; 20MG	A075776 003	Jul 01, 2002
MYLAN	12.5MG; 10MG	A076113 001	Jul 01, 2002
	12.5MG; 20MG	A076113 002	Jul 01, 2002
	25MG; 20MG	A076113 003	Jul 01, 2002
SANDOZ	12.5MG; 10MG	A075926 001	Jul 01, 2002
	12.5MG; 20MG	A075926 002	Jul 01, 2002
	25MG; 20MG	A075926 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

HYZAAR

+ MERCK SHARP DOHME	12.5MG; 50MG	N020387 001	Apr 28, 1995
+	25MG; 100MG	N020387 002	Nov 10, 1998

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
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

HYDROCHLOROTHIAZIDE; METHYLDOPA

TABLET; ORAL

ALDORIL 15

MERCK	15MG; 250MG	N013402 001	
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ALDORIL 25

MERCK	25MG; 250MG	N013402 002	
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ALDORIL D30

MERCK	30MG; 500MG	N013402 003	
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ALDORIL D50

MERCK	50MG; 500MG	N013402 004	
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METHYLDOPA AND HYDROCHLOROTHIAZIDE

DAVA PHARMS INC	15MG; 250MG	A072507 001	Jun 02, 1989
	25MG; 250MG	A072508 001	Jun 02, 1989

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

HYDROCHLOROTHIAZIDE; METHYLDOPA

TABLET; ORAL

METHYLDOPA AND HYDROCHLOROTHIAZIDE

	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
PAR PHARM	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
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
SANDOZ	15MG; 250MG	A070829	001	Mar 09, 1987
	25MG; 250MG	A070830	001	Mar 09, 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
YAOPHARMA CO LTD	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

HYDROCHLOROTHIAZIDE; METOPROLOL TARTRATE

TABLET; ORAL

LOPRESSOR HCT

+ US PHARMS HOLDINGS I 50MG; 100MG ** N018303 003 Dec 31, 1984

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
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

ACCORD HLTHCARE	12.5MG; 20MG	A209281	001	Feb 07, 2019
	12.5MG; 40MG	A209281	002	Feb 07, 2019
	25MG; 40MG	A209281	003	Feb 07, 2019
MYLAN	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
	25MG; 40MG	A200532	003	Apr 24, 2017

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

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
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INDERIDE LA 160/50

WYETH AYERST	50MG;160MG	N019059 003	Jul 03, 1985
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INDERIDE LA 80/50

WYETH AYERST	50MG;80MG	N019059 001	Jul 03, 1985
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TABLET; ORAL

INDERIDE-40/25

+ WYETH PHARMS INC	25MG;40MG **	N018031 001	
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INDERIDE-80/25

+ WYETH PHARMS INC	25MG;80MG **	N018031 002	
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PROPRANOLOL HYDROCHLORIDE & HYDROCHLOROTHIAZIDE

DURAMED PHARMS BARR	25MG;40MG	A071126 001	Mar 02, 1987
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	25MG;80MG	A071127 001	Mar 02, 1987
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PROPRANOLOL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE

ACTAVIS ELIZABETH	25MG;40MG	A070851 001	May 15, 1986
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	25MG;80MG	A070852 001	May 15, 1986
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ANI PHARMS INC	25MG;40MG	A070704 001	Oct 01, 1986
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	25MG;40MG	A072042 001	Mar 14, 1988
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	25MG;80MG	A070705 001	Oct 01, 1986
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	25MG;80MG	A072043 001	Mar 14, 1988
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IVAX SUB TEVA PHARMS	25MG;40MG	A071552 001	Dec 01, 1988
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	25MG;80MG	A071553 001	Dec 01, 1988
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WARNER CHILCOTT	25MG;40MG	A071771 001	Jan 26, 1988
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	25MG;80MG	A071772 001	Jan 26, 1988
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WATSON LABS	25MG;40MG	A070301 001	Apr 18, 1986
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	25MG;40MG	A071498 001	Dec 18, 1991
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	25MG;80MG	A070305 001	Apr 18, 1986
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	25MG;80MG	A071501 001	Dec 18, 1991
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YAOPHARMA CO LTD	25MG;40MG	A071060 001	Aug 26, 1987
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	25MG;80MG	A071061 001	Aug 26, 1987
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HYDROCHLOROTHIAZIDE; QUINAPRIL HYDROCHLORIDE

TABLET; ORAL

QUINAPRIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE

MYLAN	12.5MG;EQ 10MG BASE	A077093 001	Mar 28, 2005
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	12.5MG;EQ 20MG BASE	A077093 002	Mar 28, 2005
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	25MG;EQ 20MG BASE	A077093 003	Mar 28, 2005
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SUN PHARM INDS LTD	12.5MG;EQ 10MG BASE	A078211 001	Mar 04, 2009
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	12.5MG;EQ 20MG BASE	A078211 002	Mar 04, 2009
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	25MG;EQ 20MG BASE	A078211 003	Mar 04, 2009
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HYDROCHLOROTHIAZIDE; RESERPINE

TABLET; ORAL

H.R.-50

WHITEWORTH TOWN PLSN	50MG;0.125MG	A085338 001	
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HYDRO-RESERP

ABC HOLDING	50MG;0.125MG	A084714 002	Jun 29, 1982
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HYDRO-SERP "25"

SANDOZ	25MG;0.125MG	A084827 001	
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HYDRO-SERP "50"

SANDOZ	50MG;0.125MG	A085213 001	
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HYDROCHLOROTHIAZIDE W/ RESERPINE

IVAX SUB TEVA PHARMS	25MG;0.1MG	A083572 001	
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	25MG;0.125MG	A083571 001	
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	50MG;0.1MG	A083568 001	
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	50MG;0.125MG	A083573 001	
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PHARMERAL	25MG;0.125MG	A085421 001	
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	50MG;0.125MG	A085420 001	
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ROXANE	50MG;0.125MG	A084603 001	
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WATSON LABS	25MG;0.125MG	A084466 001	
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	25MG;0.125MG	A085317 001	
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	25MG;0.125MG	A086330 002	
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Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

HYDROCHLOROTHIAZIDE; RESERPINE

TABLET; ORAL

HYDROCHLOROTHIAZIDE W/ RESERPINE

	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	

HYDROCHLOROTHIAZIDE; SPIRONOLACTONE

TABLET; ORAL

SPIRONOLACTONE AND HYDROCHLOROTHIAZIDE

ASCOT	25MG;25MG	A088025	001	Nov 23, 1984
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	
YAOPHARMA CO LTD	25MG;25MG	A086881	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; TIMOLOL MALEATE

TABLET; ORAL

TIMOLIDE 10-25

MERCK	25MG;10MG	N018061	001	
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HYDROCHLOROTHIAZIDE; TRIAMTERENE

CAPSULE; ORAL

DYAZIDE

GLAXOSMITHKLINE LLC	25MG;50MG	N016042	002	
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TRIAMTERENE AND HYDROCHLOROTHIAZIDE

ANI PHARMS INC	25MG;37.5MG	A074970	001	Jan 06, 1998
	25MG;50MG	A074259	001	Mar 30, 1995
CASI PHARMS INC	25MG;50MG	A073191	001	Jul 31, 1991
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 INC	50MG;75MG	A073467	001	Jan 31, 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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

HYDROCHLOROTHIAZIDE; VALSARTAN

TABLET; ORAL

VALSARTAN AND HYDROCHLOROTHIAZIDE

WATSON LABS TEVA	12.5MG; 80MG	A091519 001	Mar 21, 2013
	12.5MG; 160MG	A091519 002	Mar 21, 2013
	12.5MG; 320MG	A091519 003	Mar 21, 2013
	25MG; 160MG	A091519 004	Mar 21, 2013
	25MG; 320MG	A091519 005	Mar 21, 2013
ZYDUS PHARMS	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

TABLET, EXTENDED RELEASE; ORAL

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 BITARTRATE AND IBUPROFEN

ANI PHARMS INC	5MG; 200MG	A077454 001	Jun 23, 2010
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
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
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HYDROCODONE BITARTRATE; PSEUDOEPHEDRINE HYDROCHLORIDE

SOLUTION; ORAL

HYDROCODONE BITARTRATE AND PSEUDOEPHEDRINE HYDROCHLORIDE

MAYNE PHARMA INC	5MG/5ML; 60MG/5ML	A205658 001	Nov 17, 2015
PADDOCK LLC	5MG/5ML; 60MG/5ML	A204658 001	Apr 29, 2014
TORRENT	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	
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HYDROCORTISONE

AEROSOL; TOPICAL

AEROSEB-HC

ALLERGAN HERBERT	0.5%	A085805 001	
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CREAM; TOPICAL

CORT-DOME

BAYER PHARMS	0.5%	N009585 003	
	1%	N009585 001	

DERMACORT

MONARCH PHARMS	1%	A083011 002	
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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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

HYDROCORTISONE

CREAM;TOPICAL

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

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

TARO PHARM INDS LTD 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

PENECORT

ALLERGAN HERBERT 1% A088216 001 Jun 06, 1984

PROCTOCORT

MONARCH PHARMS 1% A083011 001

SYNACORT

MEDICIS 0.5% A087459 001

1% A087458 001

2.5% A087457 001

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

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

HYDROCORTISONE

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
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				
PERRIGO CO	1%	A089066	001	Nov 25, 1985
OINTMENT; TOPICAL				
CORTRIL				
PFIZER GLOBAL	1%	N009176	001	
	2.5%	N009176	002	
HC (HYDROCORTISONE)				
C AND M PHARMA	0.5%	A080481	001	
	1%	A080481	002	
HYDROCORTISONE				
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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

6-219(of 430)

** See List Footnote

HYDROCORTISONE

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

ELKINS SINN 20MG A080624 001

FERRANTE 10MG A080568 001

20MG A080568 002

IMPAX LABS 20MG A080781 001

INWOOD LABS 20MG A080732 001

LANNETT 20MG A085070 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

WATSON LABS 20MG A080355 001

WHITEWORTH TOWN PLSN 10MG A080344 001

20MG A080344 002

HYDROCORTONE

MERCK 10MG N008506 007

20MG N008506 011

TABLET; VAGINAL

CORTRIL

PFIPHARMECS 10MG N009796 001

HYDROCORTISONE ACETATE

CREAM; TOPICAL

HEMSOL-HC

ABLE 1% A081274 001 Jun 19, 1992

HYDROCORTISONE ACETATE

CENCI 1% A080419 001 Jan 25, 1982

FERNDAL LABS 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

INJECTABLE; INJECTION

CORTEF ACETATE

PHARMACIA AND UPJOHN 50MG/ML N009378 002

CORTRIL

PFIZER 25MG/ML N009164 001

HYDROCORTISONE ACETATE

AKORN 25MG/ML N009637 001

50MG/ML N009637 002

BEL MAR 25MG/ML A083739 001

50MG/ML A083739 002

WATSON LABS 25MG/ML A083128 001

25MG/ML A083759 001

50MG/ML A083759 002

50MG/ML A085214 001

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

6-220(of 430)

** See List Footnote

HYDROCORTISONE ACETATE

INJECTABLE; INJECTION

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

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

AKORN

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; 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 ACETATE; UREA

CREAM; TOPICAL

CARMOL HC

FOUGERA PHARMS

1%;10%

A080505 001

HYDROCORTISONE BUTYRATE

CREAM; TOPICAL

LOCOID

YAMANOUCHI

0.1%

N018795 001 Jan 07, 1983

OINTMENT; TOPICAL

LOCOID

YAMANOUCHI

0.1%

N019106 001 Jul 03, 1984

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

6-221(of 430)

** See List Footnote

HYDROCORTISONE BUTYRATESOLUTION; TOPICAL
LOCOID

YAMAOUCHI 0.1% N019819 001 Sep 15, 1988

HYDROCORTISONE CYPIONATESUSPENSION; ORAL
CORTEF

PHARMACIA AND UPJOHN EQ 10MG BASE/5ML N009900 001

HYDROCORTISONE SODIUM PHOSPHATEINJECTABLE; INJECTION
HYDROCORTONE

MERCK EQ 50MG BASE/ML N012052 001

HYDROCORTISONE SODIUM SUCCINATEINJECTABLE; 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
 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

ACP NIMBLE 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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

6-222(of 430)

** See List Footnote

HYDROCORTISONE; NEOMYCIN SULFATE; POLYMYXIN B SULFATE

SOLUTION/DROPS;OTIC

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

HYDROCORTISONE; POLYMYXIN B SULFATE

SOLUTION/DROPS;OTIC

OTOBIO TIC

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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

6-223(of 430)

** See List Footnote

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

+ FRESENIUS KABI USA	4MG/ML	N019034 005	Apr 30, 2009
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DILAUDID-HP

+ FRESENIUS KABI USA	10MG/ML	N019034 001	Jan 11, 1984
	250MG/VIAL	N019034 002	Aug 04, 1994

HYDROMORPHONE HYDROCHLORIDE

HOSPIRA	10MG/ML	A074598 001	Jun 19, 1997
WATSON LABS	10MG/ML	A074317 001	Aug 23, 1995

TABLET;ORAL

HYDROMORPHONE HYDROCHLORIDE

NESHER PHARMS	2MG	A077311 001	Nov 09, 2005
	4MG	A077311 002	Nov 09, 2005
	8MG	A077311 003	Nov 09, 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	
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CYANOKIT

SERB SA	2.5GM/VIAL (5GM/KIT)	N022041 002	Dec 15, 2006
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HYDROXOCOBALAMIN

ABRAXIS PHARM	1MG/ML	A084921 001	
WATSON LABS	1MG/ML	A085528 001	

HYDROXOMIN

BEL MAR	1MG/ML	A084629 001	
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HYDROXYAMPHETAMINE HYDROBROMIDE

SOLUTION/DROPS;OPHTHALMIC

PAREDRIINE

PHARMICS	1%	N000004 004	
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HYDROXYCHLOROQUINE SULFATE

TABLET;ORAL

HYDROXYCHLOROQUINE SULFATE

INVATECH	200MG	A040150 001	Jan 27, 1996
LAURUS LABS LTD	200MG	A210959 001	Jan 15, 2019
LUPIN LTD	200MG	A210543 001	Jul 06, 2018

HYDROXYPROGESTERONE CAPROATE

INJECTABLE;INJECTION

HYDROXYPROGESTERONE CAPROATE

AKORN	125MG/ML	N018004 001	
ALLERGAN	125MG/ML	N017439 001	
	250MG/ML	N017439 002	

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

AM REGENT	1250MG/5ML (250MG/ML)	A210724 001	Aug 09, 2019
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Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

6-224(of 430)

** See List Footnote

HYDROXYPROGESTERONE CAPROATE

SOLUTION; INTRAMUSCULAR

HYDROXYPROGESTERONE CAPROATE

EUGIA PHARMA 1250MG/5ML (250MG/ML) A211142 001 May 09, 2019

HYDROXYSTILBAMIDINE ISETHIONATE

INJECTABLE; INJECTION

HYDROXYSTILBAMIDINE ISETHIONATE

SANOFI AVENTIS US 225MG/AMP N009166 001

HYDROXYUREA

CAPSULE; ORAL

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

KV PHARM 10MG/5ML A087730 001 Jul 01, 1982

TORRENT 10MG/5ML A210634 001 Feb 26, 2019

TABLET; ORAL

ATARAX

+ PFIZER 10MG ** N010392 001

+ 25MG ** N010392 004

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

6-225(of 430)

** 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

AUROLIFE PHARMA LLC

10MG

A087871 002 Dec 20, 1982

25MG

A087871 003 Dec 20, 1982

50MG

A087871 001 Dec 20, 1982

HALSEY

10MG

A089366 001 May 02, 1988

25MG

A089117 001 May 02, 1988

50MG

A089396 001 May 02, 1988

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

MUTUAL PHARM

10MG

A088409 001 Nov 15, 1983

25MG

A087857 001 Apr 18, 1983

50MG

A087860 001 Apr 18, 1983

MYLAN

10MG

A091176 001 Jun 07, 2010

25MG

A091176 002 Jun 07, 2010

50MG

A091176 003 Jun 07, 2010

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

SANDOZ

10MG

A087246 002

25MG

A085247 001

50MG

A087245 001

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

SUPERPHARM

100MG

A087862 001 Apr 18, 1983

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

VINTAGE

10MG

A087602 001 Jan 22, 1982

25MG

A087603 001 Jan 22, 1982

50MG

A087604 001 Jan 22, 1982

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

DISCONTINUED DRUG PRODUCT LIST

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** 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 25MG HYDROCHLORIDE N011459 002

+ EQ 50MG HYDROCHLORIDE N011459 004

EQ 100MG HYDROCHLORIDE ** N011459 006

SUSPENSION; ORAL

VISTARIL

PFIZER EQ 25MG HYDROCHLORIDE/5ML N011795 001

IBANDRONATE SODIUM

INJECTABLE; INTRAVENOUS

IBANDRONATE SODIUM

EMCURE PHARMS LTD EQ 3MG BASE/3ML A203987 001 Sep 02, 2014

TABLET; ORAL

BONIVA

+ HOFFMANN LA ROCHE EQ 2.5MG BASE ** N021455 001 May 16, 2003

IBANDRONATE SODIUM

MYLAN PHARMS INC EQ 150MG BASE A078995 001 Mar 19, 2012

SUN PHARM INDUSTRIES EQ 150MG BASE A078996 001 Aug 15, 2012

IBUPROFEN

CAPSULE; ORAL

IBUPROFEN

CONTRACT PHARMACAL 200MG A074782 001 Jul 06, 1998

STRIDES PHARMA EQ 200MG FREE ACID AND POTASSIUM SALT A204469 001 Mar 28, 2018

MIDOL

BAYER 200MG ** A070626 001 Sep 02, 1987

200MG ** A071002 001 Sep 02, 1987

SOLUTION; INTRAVENOUS

CALDOLOR

CUMBERLAND PHARMS 400MG/4ML (100MG/ML) N022348 001 Jun 11, 2009

SUSPENSION; ORAL

CHILDREN'S ADVIL

GLAXOSMITHKLINE 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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

IBUPROFEN

SUSPENSION;ORAL

IBUPROFEN

GUARDIAN DRUG 100MG/5ML A210149 001 Aug 17, 2018

MOTRIN

+ MCNEIL CONSUMER 100MG/5ML ** N019842 001 Sep 19, 1989

SUSPENSION/DROPS;ORAL

MOTRIN

MCNEIL 40MG/ML N020476 001 May 25, 1995

PEDIATRIC ADVIL

+ GLAXOSMITHKLINE 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 800MG A071965 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

ANI PHARMS INC 200MG A071144 001 Jan 20, 1987

200MG A072901 001 Dec 19, 1991

200MG A072903 001 Dec 19, 1991

AUROLIFE PHARMA LLC 300MG A070736 002 Jun 12, 1986

400MG A070736 003 Jun 12, 1986

600MG A070736 001 Jun 12, 1986

800MG A071938 001 Jan 14, 1988

CONTRACT PHARMACAL 200MG A071265 001 Oct 15, 1986

200MG A071265 002 Sep 10, 1987

200MG A071735 001 Sep 10, 1987

200MG A073691 001 Feb 25, 1994

200MG A074931 001 Jul 20, 1998

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

HEC PHARM 400MG A204062 001 Sep 10, 2018

600MG A204062 002 Sep 10, 2018

800MG A204062 003 Sep 10, 2018

IVAX SUB TEVA PHARMS 200MG A071154 001 Oct 27, 1987

200MG A072040 001 Apr 29, 1988

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

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

MYLAN 200MG A071870 001 May 05, 1988

400MG A070045 001 Sep 24, 1985

600MG A070057 001 Sep 24, 1985

800MG A071999 001 Dec 03, 1987

NORTHSTAR HLTHCARE 400MG A078132 001 Sep 10, 2007

600MG A078132 002 Sep 10, 2007

800MG A078132 003 Sep 10, 2007

OHM LABS 400MG A070818 001 Dec 26, 1985

P AND L DEV LLC 200MG A070733 001 Sep 19, 1986

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

IBUPROFEN

TABLET; ORAL

IBUPROFEN

PAR PHARM	200MG		A071575	001	May 08, 1987
	300MG		A070328	001	Aug 06, 1985
	400MG		A070329	001	Aug 06, 1985
	600MG		A070330	001	Aug 06, 1985
	800MG		A070986	001	Jul 25, 1986
PERRIGO	200MG		A072098	001	Dec 08, 1987
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
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
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
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
VINTAGE PHARMS	200MG		A072249	001	Jan 10, 1989
	300MG		A071230	001	Oct 22, 1986
	400MG		A071231	001	Oct 22, 1986
	600MG		A071232	001	Oct 22, 1986
	800MG		A072004	001	Nov 18, 1987
WATSON LABS	200MG		A070435	001	Mar 05, 1986
	200MG		A071765	001	Sep 04, 1987
	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
IBUPROHM					
OHM LABS	200MG		A071214	001	Dec 01, 1986
	400MG		A070469	001	Aug 29, 1985
MEDIPREN					
MCNEIL	200MG		A070475	001	Feb 06, 1986
	200MG		A071215	001	Jun 26, 1986
MIDOL					
BAYER	200MG		A070591	001	Sep 02, 1987
	200MG		A071001	001	Sep 02, 1987
MOTRIN					
+ MCNEIL CONSUMER	300MG **		N017463	003	
+	400MG **		N017463	002	
+	600MG **		N017463	004	
+	800MG **		N017463	005	May 22, 1985
MCNEIL PED	100MG		N020418	001	Nov 16, 1994
MOTRIN MIGRAINE PAIN					
J AND J CONSUMER INC	200MG		N019012	004	Feb 25, 2000
NUPRIN					
BRISTOL MYERS	200MG		A072035	001	Feb 16, 1988

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

IBUPROFENTABLET; ORAL
NUPRIN

	200MG	A072036 001	Feb 16, 1988
J AND J CONSUMER INC	200MG	N019012 001	May 18, 1984
	200MG	N019012 002	Jul 29, 1987
RUFEN			
BASF	600MG	N018197 002	Mar 05, 1984
TABLET, CHEWABLE; ORAL			
CHILDREN'S MOTRIN			
+ J AND J CONSUMER INC	50MG	N020601 001	Nov 15, 1996
IBUPROFEN			
PERRIGO	50MG	A076359 001	Jan 16, 2004
JUNIOR STRENGTH MOTRIN			
+ J AND J CONSUMER INC	100MG	N020601 003	Nov 15, 1996
MOTRIN			
MCNEIL PED	50MG	N020135 001	Nov 16, 1994
	100MG	N020135 002	Nov 16, 1994

IBUPROFEN; OXYCODONE HYDROCHLORIDE

TABLET; ORAL

COMBUNOX			
FOREST LABS	400MG; 5MG **	N021378 001	Nov 26, 2004
OXYCODONE HYDROCHLORIDE AND IBUPROFEN			
WATSON LABS	400MG; 5MG	A078394 001	Nov 26, 2007

IBUPROFEN; PSEUDOEPHEDRINE HYDROCHLORIDE

TABLET; ORAL

IBUPROFEN AND PSEUDOEPHEDRINE HYDROCHLORIDE			
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

IDARUBICIN HYDROCHLORIDE

INJECTABLE; INJECTION

IDAMYCIN			
PHARMACIA AND UPJOHN	5MG/VIAL	N050661 002	Sep 27, 1990
	10MG/VIAL	N050661 001	Sep 27, 1990
	20MG/VIAL	N050661 003	Apr 25, 1995
IDARUBICIN HYDROCHLORIDE			
MYLAN LABS LTD	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			
FRESENIUS KABI USA	1GM/20ML (50MG/ML)	A090181 001	Sep 22, 2009
	3GM/60ML (50MG/ML)	A090181 002	Sep 22, 2009
MYLAN LABS LTD	1GM/20ML (50MG/ML)	A201689 001	Nov 26, 2012
	3GM/60ML (50MG/ML)	A201689 002	Nov 26, 2012

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

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

10MG

A207231 006 Nov 28, 2016

12MG

A207231 007 Nov 28, 2016

TARO PHARM INDS LTD

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 PHARMS LTD

20MCG/2ML (10MCG/ML)

N021779 001 Dec 29, 2004

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

IMIPRAMINE HYDROCHLORIDE

CONCENTRATE; ORAL

IMIPRAMINE HYDROCHLORIDE

NOVARTIS

25MG/ML

A086765 001

INJECTABLE; INJECTION

TOFRANIL

NOVARTIS

12.5MG/ML

N011838 002

TABLET; ORAL

IMIPRAMINE HYDROCHLORIDE

LEDERLE

10MG

A086269 001

25MG

A086267 001

50MG

A086268 001

LUPIN LTD

10MG

A090441 002 Mar 11, 2010

25MG

A090441 003 Mar 11, 2010

50MG

A090441 001 Mar 11, 2010

OXFORD PHARMS

10MG

A040751 003 Feb 28, 2008

25MG

A040751 002 Feb 28, 2008

50MG

A040751 001 Feb 28, 2008

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

TEVA

10MG

A083729 001

25MG

A083729 004

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

IMIPRAMINE HYDROCHLORIDE

TABLET; ORAL

IMIPRAMINE HYDROCHLORIDE

	50MG	A083729 003	
USL PHARMA	25MG	A087776 001	Feb 10, 1982
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	

IMIPRAMINE PAMOATE

CAPSULE; ORAL

IMIPRAMINE PAMOATE

MYLAN PHARMS INC	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

IMIQUIMOD

ACP NIMBLE	5%	A200481 001	Apr 18, 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

INDAPAMIDE

TABLET; ORAL

INDAPAMIDE

ANI PHARMS INC	1.25MG	A074498 002	Feb 12, 1998
	2.5MG	A074498 001	Oct 31, 1996
MYLAN PHARMS INC	1.25MG	A075105 001	Jul 23, 1998
	2.5MG	A075105 002	Jul 23, 1998
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
YAOPHARMA CO LTD	1.25MG	A074594 001	May 23, 1996
	2.5MG	A074594 002	May 23, 1996
LOZOL			
+ SANOFI AVENTIS US	1.25MG **	N018538 002	Apr 29, 1993

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

INDAPAMIDE

TABLET; ORAL

LOZOL

+

2.5MG **

N018538 001 Jul 06, 1983

INDECAINIDE HYDROCHLORIDE

TABLET, 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 333MG BASE

N020685 005 Dec 17, 1998

INDIUM IN-111 CHLORIDE

INJECTABLE; INJECTION

INDICLOR

+

GE HEALTHCARE

2mCi/0.2ML

N019862 001 Dec 29, 1992

INDOCYANINE GREEN

INJECTABLE; INJECTION

IC-GREEN

AKORN

10MG/VIAL **

N011525 003

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

+

EGALET

25MG **

N016059 001

+

50MG **

N016059 002

INDOMETHACIN

ABLE

25MG

A076666 001 Dec 17, 2003

50MG

A076666 002 Dec 17, 2003

ANI PHARMS INC

25MG

A071148 001 Mar 18, 1987

50MG

A071149 001 Mar 18, 1987

CHARTWELL MOLECULES

25MG

N018829 002 Aug 06, 1984

50MG

A070651 001 Mar 05, 1986

50MG

N018829 001 Aug 06, 1984

CYCLE PHARMS LTD

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

IVAX SUB TEVA PHARMS

25MG

N018730 001 May 04, 1984

50MG

N018730 002 May 04, 1984

MUTUAL PHARM

25MG

A070067 001 Oct 03, 1986

50MG

A070068 001 Oct 03, 1986

MYLAN

25MG

N018858 001 Apr 20, 1984

50MG

A070624 001 Sep 04, 1985

50MG

N018858 002 Apr 20, 1984

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

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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

INDOMETHACIN

CAPSULE; ORAL

INDOMETHACIN

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

CAPSULE, EXTENDED RELEASE; ORAL

INDOCIN SR

+ EGALET

75MG **

N018185 001 Feb 23, 1982

INDOMETHACIN

ABLE

75MG

A076114 001 Feb 06, 2002

INWOOD LABS

75MG

A072410 001 Mar 15, 1989

WATSON LABS INC

75MG

A202572 001 Dec 09, 2013

SUPPOSITORY; RECTAL

INDOCIN

+ EGALET

50MG **

N017814 001 Aug 13, 1984

SUSPENSION; ORAL

INDOMETHACIN

HIKMA

25MG/5ML

A071412 001 Mar 18, 1987

INSULIN ASPART PROTAMINE RECOMBINANT; INSULIN ASPART RECOMBINANT

INJECTABLE; SUBCUTANEOUS

NOVOLOG MIX 50/50

NOVO NORDISK INC

50 UNITS/ML; 50 UNITS/ML

N021810 001 Aug 26, 2008

NOVOLOG MIX 70/30 PENFILL

NOVO NORDISK INC

210 UNITS/3ML; 90 UNITS/3ML (70 UNITS/ML; 30 UNITS/ML)

N021172 002 Nov 01, 2001

210 UNITS/3ML; 90 UNITS/3ML (70 UNITS/ML; 30 UNITS/ML)

N021172 003 Nov 01, 2001

INSULIN ASPART RECOMBINANT

INJECTABLE; SUBCUTANEOUS

NOVOLOG FLEXTOUCH

+ NOVO NORDISK INC

300 UNITS/3ML (100 UNITS/ML)

N020986 005 Oct 31, 2013

NOVOLOG INNOLET

NOVO NORDISK INC

300 UNITS/3ML (100 UNITS/ML)

N020986 004 Apr 23, 2004

INSULIN ASPART; INSULIN DEGLUDEC

SOLUTION; SUBCUTANEOUS

RYZODEG 70/30

+ NOVO

90 UNITS/3ML; 210 UNITS/3ML (30 UNITS/ML; 70 UNITS/ML)

N203313 001 Sep 25, 2015

INSULIN DETEMIR RECOMBINANT

INJECTABLE; SUBCUTANEOUS

LEVEMIR FLEXPEN

NOVO NORDISK INC

300 UNITS/3ML (100 UNITS/ML)

N021536 002 Jun 16, 2005

LEVEMIR INNOLET

NOVO NORDISK INC

300 UNITS/3ML (100 UNITS/ML)

N021536 003 Jun 16, 2005

LEVEMIR PENFILL

NOVO NORDISK INC

300 UNITS/3ML (100 UNITS/ML)

N021536 004 Jun 16, 2005

INSULIN GLULISINE RECOMBINANT

INJECTABLE; INTRAVENOUS, SUBCUTANEOUS

APIDRA

+ SANOFI AVENTIS US

300 UNITS/3ML (100 UNITS/ML)

N021629 002 Dec 20, 2005

INSULIN LISPRO PROTAMINE RECOMBINANT; INSULIN LISPRO RECOMBINANT

INJECTABLE; INJECTION

HUMALOG MIX 50/50 PEN

LILLY

50 UNITS/ML; 50 UNITS/ML

N021018 003 Dec 22, 1999

HUMALOG MIX 75/25 PEN

LILLY

75 UNITS/ML; 25 UNITS/ML

N021017 003 Dec 22, 1999

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

INSULIN LISPRO RECOMBINANT

INJECTABLE; INJECTION

HUMALOG PEN

LILLY

100 UNITS/ML

N020563 002 Aug 06, 1998

INSULIN PORK

INJECTABLE; INJECTION

ILETIN I

LILLY

500 UNITS/ML

N017931 001

INSULIN

NOVO NORDISK INC

40 UNITS/ML

N017926 001

REGULAR INSULIN

NOVO NORDISK INC

100 UNITS/ML

N017926 003

INSULIN PURIFIED BEEF

INJECTABLE; INJECTION

REGULAR ILETIN II

LILLY

100 UNITS/ML

N018478 001

INSULIN PURIFIED PORK

INJECTABLE; INJECTION

ILETIN II

LILLY

500 UNITS/ML

N018344 002

REGULAR ILETIN II (PORK)

LILLY

100 UNITS/ML

N018344 001

REGULAR PURIFIED PORK INSULIN

NOVO NORDISK INC

100 UNITS/ML

N018381 001

VELOSULIN

NOVO NORDISK INC

100 UNITS/ML

N018193 001

INSULIN PURIFIED PORK; INSULIN SUSP ISOPHANE PURIFIED PORK

INJECTABLE; INJECTION

INSULIN NORDISK MIXTARD (PORK)

NOVO NORDISK INC

30 UNITS/ML; 70 UNITS/ML

N018195 001

INSULIN RECOMBINANT HUMAN

INJECTABLE; INJECTION

HUMULIN BR

LILLY

100 UNITS/ML

N019529 001 Apr 28, 1986

VELOSULIN BR

NOVO NORDISK INC

100 UNITS/ML

N021028 001 Jul 19, 1999

POWDER; INHALATION

EXUBERA

PFIZER

1MG/INH

N021868 001 Jan 27, 2006

3MG/INH

N021868 002 Jan 27, 2006

INSULIN RECOMBINANT HUMAN; INSULIN SUSP ISOPHANE RECOMBINANT HUMAN

INJECTABLE; INJECTION

HUMULIN 50/50

LILLY

50 UNITS/ML; 50 UNITS/ML

N020100 001 Apr 29, 1992

INSULIN RECOMBINANT PURIFIED HUMAN

INJECTABLE; INJECTION

NOVOLIN R

NOVO NORDISK INC

100 UNITS/ML

N018778 001 Aug 30, 1983

VELOSULIN BR HUMAN

NOVO NORDISK INC

100 UNITS/ML

N019450 001 May 30, 1986

INSULIN RECOMBINANT PURIFIED HUMAN; INSULIN SUSP ISOPHANE SEMISYNTHETIC PURIFIED HUMAN

INJECTABLE; INJECTION

MIXTARD HUMAN 70/30

BAYER PHARMS

30 UNITS/ML; 70 UNITS/ML

N019585 001 Mar 11, 1988

NOVOLIN 70/30

NOVO NORDISK INC

30 UNITS/ML; 70 UNITS/ML

N019441 001 Jul 11, 1986

INSULIN SUSP ISOPHANE BEEF

INJECTABLE; INJECTION

NPH INSULIN

NOVO NORDISK INC

40 UNITS/ML

N017929 001

100 UNITS/ML

N017929 003

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

INSULIN SUSP ISOPHANE BEEF/PORK

INJECTABLE; INJECTION

NPH ILETIN I (BEEF-PORK)

LILLY

40 UNITS/ML

N017936 001

100 UNITS/ML

N017936 002

INSULIN SUSP ISOPHANE PURIFIED BEEF

INJECTABLE; INJECTION

NPH ILETIN II

LILLY

100 UNITS/ML

N018479 001

INSULIN SUSP ISOPHANE PURIFIED PORK

INJECTABLE; INJECTION

INSULIN INSULATARD NPH NORDISK

NOVO NORDISK INC

100 UNITS/ML

N018194 001

NPH ILETIN II (PORK)

LILLY

100 UNITS/ML

N018345 001

NPH PURIFIED PORK ISOPHANE INSULIN

NOVO NORDISK INC

100 UNITS/ML

N018623 001

INSULIN SUSP ISOPHANE SEMISYNTHETIC PURIFIED HUMAN

INJECTABLE; INJECTION

INSULATARD NPH HUMAN

NOVO NORDISK INC

100 UNITS/ML

N019449 001 May 30, 1986

NOVOLIN N

NOVO NORDISK INC

100 UNITS/ML

N019065 001 Jan 23, 1985

INSULIN SUSP PROTAMINE ZINC BEEF/PORK

INJECTABLE; INJECTION

PROTAMINE ZINC & ILETIN I (BEEF-PORK)

LILLY

40 UNITS/ML

N017932 001

100 UNITS/ML

N017932 002

INSULIN SUSP PROTAMINE ZINC PURIFIED BEEF

INJECTABLE; INJECTION

PROTAMINE ZINC AND ILETIN II

LILLY

100 UNITS/ML

N018476 001

PROTAMINE ZINC INSULIN

BRISTOL MYERS SQUIBB

40 UNITS/ML

N017928 001

100 UNITS/ML

N017928 003

INSULIN SUSP PROTAMINE ZINC PURIFIED PORK

INJECTABLE; INJECTION

PROTAMINE ZINC AND ILETIN II (PORK)

LILLY

100 UNITS/ML

N018346 001

INSULIN ZINC SUSP BEEF

INJECTABLE; INJECTION

LENTE INSULIN

NOVO NORDISK INC

40 UNITS/ML

N017998 001

100 UNITS/ML

N017998 003

INSULIN ZINC SUSP EXTENDED BEEF

INJECTABLE; INJECTION

ULTRALENTE INSULIN

NOVO NORDISK INC

100 UNITS/ML

N017997 003

INSULIN ZINC SUSP EXTENDED PURIFIED BEEF

INJECTABLE; INJECTION

ULTRALENTE

NOVO NORDISK INC

100 UNITS/ML

N018385 001

INSULIN ZINC SUSP EXTENDED RECOMBINANT HUMAN

INJECTABLE; INJECTION

HUMULIN U

LILLY

40 UNITS/ML

N019571 001 Jun 10, 1987

100 UNITS/ML

N019571 002 Jun 10, 1987

INSULIN ZINC SUSP PROMPT BEEF

INJECTABLE; INJECTION

SEMILENTE INSULIN

NOVO NORDISK INC

100 UNITS/ML

N017996 003

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

INSULIN ZINC SUSP PROMPT PURIFIED PORKINJECTABLE; INJECTION
SEMILENTE

NOVO NORDISK INC 100 UNITS/ML N018382 001

INSULIN ZINC SUSP PURIFIED BEEFINJECTABLE; INJECTION
LENTE ILETIN II

LILLY 100 UNITS/ML N018477 001

INSULIN ZINC SUSP PURIFIED BEEF/PORKINJECTABLE; INJECTION
LENTARD

NOVO NORDISK INC 100 UNITS/ML N018384 001

INSULIN ZINC SUSP PURIFIED PORKINJECTABLE; INJECTION
LENTE

NOVO NORDISK INC 100 UNITS/ML N018383 001

LENTE ILETIN II (PORK)
LILLY 100 UNITS/ML N018347 001INSULIN ZINC SUSP RECOMBINANT HUMANINJECTABLE; INJECTION
HUMULIN L

LILLY 100 UNITS/ML N019377 002 Sep 30, 1985

NOVOLIN L
NOVO NORDISK INC 100 UNITS/ML N019965 001 Jun 25, 1991INSULIN ZINC SUSP SEMISYNTHETIC PURIFIED HUMANINJECTABLE; INJECTION
NOVOLIN L

NOVO NORDISK INC 100 UNITS/ML N018777 001 Aug 30, 1983

INULININJECTABLE; INJECTION
INULIN AND SODIUM CHLORIDE

ISO TEX 100MG/ML N002282 001

INVERT SUGARINJECTABLE; INJECTION
TRAVERT 10% IN PLASTIC CONTAINER

BAXTER HLTHCARE 10GM/100ML N016717 001

IOBENGUANE SULFATE I-131INJECTABLE; INJECTION
IOBENGUANE SULFATE I 131

PHARMALUCENCE 2.3mCi/ML N020084 001 Mar 25, 1994

IO CETAMIC ACIDTABLET; ORAL
CHOLEBRINE

MALLINCKRODT 750MG N017129 001

IODAMIDE MEGLUMINEINJECTABLE; INJECTION
RENOVUE-65

BRACCO 65% N017902 001

RENOVUE-DIP
BRACCO 24% N017903 001IODIPAMIDE MEGLUMINEINJECTABLE; INJECTION
CHOLOGRAFIN MEGLUMINE

BRACCO 10.3% N009321 007

+ 52% N009321 003

IODIPAMIDE SODIUMINJECTABLE; INJECTION
CHOLOGRAFIN SODIUM

BRACCO 20% N009321 001

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

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			
HIPPURAN 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	

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

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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

IOPAMIDOL

INJECTABLE; INJECTION

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
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ISOVUE-128

BRACCO	26%	N018735	005	Oct 21, 1986
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ISOVUE-200

BRACCO	41%	N020327	001	Oct 12, 1994
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IOPANOIC ACID

TABLET; ORAL

TELEPAQUE

GE HEALTHCARE	500MG	N008032	001	
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IOPHENDYLATE

INJECTABLE; INJECTION

PANTOPAQUE

ALCON	100%	N005319	001	
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IOPROMIDE

INJECTABLE; INJECTION

ULTRAVIST (PHARMACY BULK)

+ BAYER HLTHCARE	49.9%	N021425	003	Mar 12, 2004
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ULTRAVIST 150

+ BAYER HLTHCARE	31.2%	N020220	004	May 10, 1995
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ULTRAVIST 240

+ BAYER HLTHCARE	49.9%	N020220	003	May 10, 1995
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ULTRAVIST 300 IN PLASTIC CONTAINER

+ BAYER HLTHCARE	62.3%	N020220	005	Nov 18, 2008
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IOTHALAMATE MEGLUMINE

INJECTABLE; INJECTION

CONRAY 30

+ LIEBEL-FLARSHEIM	30%	N016983	001	
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CONRAY 43

+ LIEBEL-FLARSHEIM	43%	N013295	002	
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IOTHALAMATE MEGLUMINE; IOTHALAMATE SODIUM

INJECTABLE; INJECTION

VASCORAY

MALLINCKRODT	52%;26%	N016783	001	
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IOTHALAMATE SODIUM

INJECTABLE; INJECTION

ANGIO-CONRAY

MALLINCKRODT	80%	N013319	001	
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CONRAY 325

MALLINCKRODT	54.3%	N017685	001	
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CONRAY 400

MALLINCKRODT	66.8%	N014295	001	
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IOTROLAN

INJECTABLE; INTRATHECAL

OSMOVIST 190

BAYER HLTHCARE	40.6%	N019580	001	Dec 07, 1989
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OSMOVIST 240

BAYER HLTHCARE	51.3%	N019580	002	Dec 07, 1989
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IOVERSOL

INJECTABLE; INJECTION

OPTIRAY 160

LIEBEL-FLARSHEIM	34%	N019710	003	Dec 30, 1988
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OPTIRAY 240

LIEBEL-FLARSHEIM	51%	N020923	001	May 28, 1998
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DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

IOXAGLATE MEGLUMINE; IOXAGLATE SODIUM

INJECTABLE; INJECTION

HEXABRIX

GUERBET	39.3%;19.6%	N018905 002	Jul 26, 1985
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IOXILAN

INJECTABLE; INJECTION

OXILAN-300

GUERBET	62%	N020316 001	Dec 21, 1995
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OXILAN-350

GUERBET	73%	N020316 002	Dec 21, 1995
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IPODATE CALCIUM

GRANULE; ORAL

ORAGRAFIN CALCIUM

BRACCO	3GM/PACKET	N012968 001	
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IPODATE SODIUM

CAPSULE; ORAL

BILIVIST

BAYER HLTHCARE	500MG	A087768 001	Aug 11, 1982
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ORAGRAFIN SODIUM

BRACCO	500MG	N012967 001	
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IPRATROPIUM BROMIDE

AEROSOL, METERED; INHALATION

ATROVENT

BOEHRINGER INGELHEIM	0.018MG/INH	N019085 001	Dec 29, 1986
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SOLUTION; INHALATION

ATROVENT

+ BOEHRINGER INGELHEIM	0.02% **	N020228 001	Sep 29, 1993
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IPRATROPIUM BROMIDE

ACTAVIS MID ATLANTIC	0.02%	A075111 001	Apr 22, 1999
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APOTEX INC	0.02%	A075441 001	Mar 28, 2001
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BAUSCH AND LOMB INC	0.02%	A075835 001	Oct 15, 2001
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MYLAN SPECIALITY LP	0.02%	A074755 001	Jan 10, 1997
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PHARMASCIENCE INC	0.02%	A075507 001	Jan 19, 2001
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ROXANE	0.02%	A075867 001	Jul 22, 2002
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TEVA PHARMS USA	0.02%	A075313 001	Feb 07, 2000
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SPRAY, METERED; NASAL

ATROVENT

+ BOEHRINGER INGELHEIM	0.021MG/SPRAY	N020393 001	Oct 20, 1995
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+	0.042MG/SPRAY	N020394 001	Oct 20, 1995
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IPRATROPIUM BROMIDE

MYLAN SPECIALITY LP	0.021MG/SPRAY	A075552 001	Mar 31, 2003
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	0.042MG/SPRAY	A075553 001	Mar 31, 2003
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IRBESARTAN

TABLET; ORAL

IRBESARTAN

AJANTA PHARMA LTD	75MG	A203685 001	Dec 10, 2015
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	150MG	A203685 002	Dec 10, 2015
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	300MG	A203685 003	Dec 10, 2015
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APOTEX INC	75MG	A200832 001	Oct 15, 2012
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	150MG	A200832 002	Oct 15, 2012
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	300MG	A200832 003	Oct 15, 2012
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HIKMA	75MG	A090201 001	Oct 15, 2012
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	150MG	A090201 002	Oct 15, 2012
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	300MG	A090201 003	Oct 15, 2012
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MYLAN PHARMS INC	75MG	A200461 001	Sep 27, 2012
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	150MG	A200461 002	Sep 27, 2012
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	300MG	A200461 003	Sep 27, 2012
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WATSON LABS INC	75MG	A090720 001	Oct 12, 2012
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	150MG	A090720 002	Oct 12, 2012
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	300MG	A090720 003	Oct 12, 2012
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DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

IRINOTECAN HYDROCHLORIDE

INJECTABLE; INJECTION

IRINOTECAN HYDROCHLORIDE

FRESENIUS KABI USA	40MG/2ML (20MG/ML)	A078188 001	Feb 27, 2008
	100MG/5ML (20MG/ML)	A078188 002	Feb 27, 2008
SANDOZ	40MG/2ML (20MG/ML)	A077994 001	Feb 27, 2008
	100MG/5ML (20MG/ML)	A077994 002	Feb 27, 2008
SANDOZ INC	40MG/2ML (20MG/ML)	A090137 001	Nov 12, 2009
	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

IRON DEXTRAN

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	

IRON SUCROSE

INJECTABLE; INTRAVENOUS

VENOFER

AM REGENT	EQ 65MG BASE/3.25ML (EQ 20MG BASE/ML)	N021135 005	Mar 29, 2013
	EQ 75MG BASE/3.75ML (EQ 20MG BASE/ML)	N021135 003	Mar 29, 2005

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	

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

ISOETHARINE HYDROCHLORIDE

SOLUTION; INHALATION

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

ISOETHARINE MESYLATE

AEROSOL, 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

ANDA REPOSITORY 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

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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

6-242(of 430)

** See List Footnote

ISONIAZID

TABLET; ORAL

ISONIAZID

+	SANDOZ	100MG	N008678	002	
		300MG	N008678	003	
+	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	
		300MG	A083178	001	
		300MG	A085784	001	
	WHITEWORTH TOWN PLSN	100MG	A080120	002	
LANIAZID					
	LANNETT	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; RIFAMPIN

CAPSULE; ORAL

RIFAMPIN AND ISONIAZID

	HIKMA INTL PHARMS	150MG;300MG	A065221	001	Jul 29, 2005
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ISOPROPAMIDE IODIDE

TABLET; ORAL

DARBID

	GLAXOSMITHKLINE	EQ 5MG BASE	N010744	001	
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ISOPROPYL ALCOHOL

SOLUTION; TOPICAL

ZURAGARD

+	ZUREX PHARMA	70% (10.5ML)	N210872	001	Apr 26, 2019
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ISOPROTERENOL HYDROCHLORIDE

AEROSOL, METERED; INHALATION

ISOPROTERENOL HYDROCHLORIDE

	3M	0.12MG/INH	N010375	004	
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	ALPHARMA US PHARMS	0.12MG/INH	A085904	001	
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ISUPREL

	SANOFI AVENTIS US	0.103MG/INH	N011178	001	
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DISC; INHALATION

NORISODRINE AEROTROL

	ABBOTT	0.25%	N016814	001	
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INJECTABLE; INJECTION

ISOPROTERENOL HYDROCHLORIDE

	ABRAXIS PHARM	0.2MG/ML	A083431	001	
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	BAXTER HLTHCARE	0.2MG/ML	A083486	001	
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	HOSPIRA	0.02MG/ML	A083283	001	
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		0.2MG/ML	A083346	001	
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	INTL MEDICATION	0.2MG/ML	A083724	001	
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SOLUTION; INHALATION

AEROLONE

	LILLY	0.25%	N007245	001	
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ISOPROTERENOL HYDROCHLORIDE

	ARMOUR PHARM	0.031%	A087935	001	Nov 18, 1982
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		0.062%	A087936	001	Nov 18, 1982
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	DEY	0.5%	A086764	001	Jan 04, 1982
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	PARKE DAVIS	0.25%	A085994	001	
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		0.5%	A085540	001	
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ISUPREL

	SANOFI AVENTIS US	0.5%	N006327	002	
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		1%	N006327	003	
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VAPO-ISO

	FISONS	0.5%	N016813	001	
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Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

ISOPROTERENOL HYDROCHLORIDETABLET;RECTAL, SUBLINGUAL
ISUPREL

SANOFI AVENTIS US	10MG	N006328	001
	15MG	N006328	002

ISOPROTERENOL HYDROCHLORIDE; PHENYLEPHRINE BITARTRATEAEROSOL, METERED; INHALATION
DUO-MEDIHALER

3M	0.16MG/INH;0.24MG/INH	N013296	001
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ISOPROTERENOL SULFATEAEROSOL, METERED; INHALATION
MEDIHALER-ISO

3M	0.08MG/INH	N010375	003
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POWDER; INHALATION
NORISODRINE

ABBVIE	10%	N006905	003
	25%	N006905	002

ISOSORBIDESOLUTION; ORAL
ISMOTIC

ALCON	100GM/220ML	N017063	001
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ISOSORBIDE DINITRATECAPSULE, EXTENDED RELEASE; ORAL
ISORDIL

WYETH AYERST	40MG	N012882	002	Jul 29, 1988
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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 INC	10MG	A086032	001	Jan 07, 1988
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

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
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Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

ISOSORBIDE DINITRATETABLET, CHEWABLE;ORAL
SORBITRATE

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

ISOSORBIDE MONONITRATE

TABLET;ORAL

ISMO

PROMIUS PHARMA

20MG

N019091 001 Dec 30, 1991

ISOSORBIDE MONONITRATE

ANI PHARMS INC

20MG

A075147 001 Nov 27, 1998

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

ISOSULFAN BLUE

INJECTABLE;INJECTION

ISOSULFAN BLUE

BELOTECA INC

1%

A210714 001 Jan 16, 2019

SOMERSET THERAPS LLC

1%

A210558 001 Jul 12, 2019

LYMPHAZURIN

+ COVIDIEN

1% **

N018310 001

ISOTRETINOIN

CAPSULE;ORAL

AC CUTANE

+ 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

40MG

A076041 003 Dec 24, 2002

ISRADIPINE

CAPSULE;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

MYLAN

5MG

A201067 001 Nov 27, 2015

10MG

A201067 002 Nov 27, 2015

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

ITRACONAZOLE

INJECTABLE; INJECTION

SPORANOX

JANSSEN PHARMS 10MG/ML N020966 001 Mar 30, 1999

SOLUTION; ORAL

ITRACONAZOLE

APOTEX 10MG/ML A208481 001 Aug 02, 2019

IVERMECTIN

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

WEST-WARD PHARMS INT 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
 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

CREAM; TOPICAL

NIZORAL

+ JANSSEN PHARMA 2% ** N019084 001 Dec 31, 1985

SUSPENSION; ORAL

NIZORAL

JANSSEN PHARMA 100MG/5ML A070767 001 Nov 07, 1986

TABLET; ORAL

KETOCONAZOLE

AAIPHARMA LLC 200MG A075341 001 Jul 27, 1999
 HAVIX 200MG A075912 001 Jan 10, 2002
 HERITAGE PHARMA 200MG A074971 001 Jun 15, 1999
 200MG A075362 001 Jun 15, 1999

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

KETOCONAZOLE

TABLET; ORAL

KETOCONAZOLE

SUN PHARM INDUSTRIES 200MG

A075314 001 Jun 15, 1999

TEVA 200MG

A075273 001 Jun 15, 1999

NIZORAL

+ JANSEN PHARMS 200MG **

N018533 001

KETOPROFEN

CAPSULE; ORAL

KETOPROFEN

AUROLIFE PHARMA LLC 50MG

A074024 001 Dec 29, 1995

75MG

A074024 002 Dec 29, 1995

MYLAN 50MG

A074035 002 Dec 31, 1996

75MG

A074035 003 Dec 31, 1996

TEVA 25MG

A073515 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

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

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

GLAND PHARMA LTD 15MG/ML

A076722 001 Jul 27, 2004

30MG/ML

A076722 002 Jul 27, 2004

HOSPIRA 15MG/ML

A074801 001 Jun 05, 1997

30MG/ML

A074801 002 Jun 05, 1997

LUITPOLD 15MG/ML

A078145 001 Jan 14, 2008

30MG/ML

A078145 002 Jan 14, 2008

MYLAN LABS LTD 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

SANDOZ INC 15MG/ML

A076271 001 Oct 06, 2004

SUN PHARM 15MG/ML

A078737 001 Oct 06, 2008

30MG/ML

A078737 002 Oct 06, 2008

WEST-WARD PHARMS INT 15MG/ML **

A075222 001 Apr 26, 1999

15MG/ML

A075299 001 Nov 03, 1999

30MG/ML **

A075222 002 Apr 26, 1999

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

KETOROLAC TROMETHAMINE

INJECTABLE; INJECTION

KETOROLAC TROMETHAMINE	30MG/ML **	A075228 001	Apr 26, 1999
	30MG/ML	A075299 002	Nov 03, 1999
WOCKHARDT	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			
AKORN	0.45%	A203376 001	Feb 10, 2014
AUROBINDO PHARMA LTD	0.4%	A205191 001	Nov 15, 2018
SANDOZ INC	0.4%	A078721 001	Nov 05, 2009
TABLET; ORAL			
KETOROLAC TROMETHAMINE			
CYCLE PHARMS LTD	10MG	A074790 001	Jun 26, 1997
WATSON LABS	10MG	A074955 001	Sep 19, 1997
TORADOL			
+ ROCHE PALO	10MG **	N019645 001	Dec 20, 1991

KETOTIFEN FUMARATE

SOLUTION/DROPS; OPHTHALMIC

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			
AKORN INC	5MG/ML	A075524 001	Nov 29, 1999
APOTHECON	5MG/ML	A075355 001	Nov 29, 1999
BAXTER HLTHCARE CORP	5MG/ML	A076051 001	Jul 05, 2002
HOSPIRA	5MG/ML	A075242 001	Sep 30, 1999
MYLAN ASI	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

TABLET; ORAL

LABETALOL HYDROCHLORIDE			
APOTHECON	100MG	A075223 001	Nov 20, 1998
	200MG	A075223 002	Nov 20, 1998
	300MG	A075223 003	Nov 20, 1998
HERITAGE PHARMA	300MG	A074787 003	Aug 03, 1998
TEVA	100MG	A074989 001	Sep 30, 1998
	200MG	A074989 002	Sep 30, 1998
	300MG	A074989 003	Sep 30, 1998
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			
+ CNTY LINE PHARMS	400MG **	N018716 004	Aug 01, 1984

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

LACTULOSE

SOLUTION; ORAL

CHRONULAC

+ SANOFI AVENTIS US 10GM/15ML **

N017884 001

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

MORTON GROVE 10GM/15ML

A071841 001 Sep 22, 1988

PACO 10GM/15ML

A073160 001 Aug 25, 1992

TORRENT 10GM/15ML

A207786 001 Jun 11, 2018

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

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

BAJAJ 10GM/15ML

A076645 001 Jul 28, 2003

PACO 10GM/15ML

A072029 001 Aug 25, 1992

ROXANE 10GM/15ML

A073590 001 May 29, 1992

SOLVAY 10GM/15ML

N017906 001

TORRENT 10GM/15ML

A203762 001 Mar 27, 2015

PORTALAC

SOLVAY 10GM/15ML

A072374 001 Mar 22, 1989

LAMIVUDINE

TABLET; ORAL

LAMIVUDINE

MYLAN 100MG

A204002 001 Dec 31, 2014

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

LAMIVUDINE; ZIDOVUDINE

TABLET; ORAL

LAMIVUDINE AND ZIDOVUDINE

MYLAN 150MG; 300MG

A079079 001 Aug 12, 2019

150MG; 300MG

A204005 001 Aug 28, 2014

PHARMACARE 150MG; 300MG

N022018 001 Mar 17, 2017

TEVA PHARMS 150MG; 300MG

A079081 001 May 25, 2011

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

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
CELLTRION	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
MYLAN	25MG	A077428	001	Jan 27, 2009
	100MG	A077428	002	Jan 27, 2009
	150MG	A077428	003	Jan 27, 2009
	200MG	A077428	004	Jan 27, 2009
MYLAN LABS LTD	25MG	A078443	001	Feb 11, 2009
	100MG	A078443	002	Feb 11, 2009
	150MG	A078443	003	Feb 11, 2009
	200MG	A078443	004	Feb 11, 2009
PHARMASCIENCE INC	25MG	A078310	001	Feb 04, 2009
	100MG	A078310	002	Feb 04, 2009
	150MG	A078310	003	Feb 04, 2009
	200MG	A078310	004	Feb 04, 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

TABLET, CHEWABLE; ORAL

LAMICTAL CD

GLAXOSMITHKLINE LLC	100MG	N020764	003	Aug 24, 1998
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LAMOTRIGINE

MYLAN	5MG	A076630	001	Jan 22, 2009
	25MG	A076630	002	Jan 22, 2009
SANDOZ	5MG	A078409	002	Jan 22, 2009
	25MG	A078409	003	Jan 22, 2009
TEVA	5MG	A076420	001	Jun 21, 2006
	25MG	A076420	002	Jun 21, 2006

TABLET, EXTENDED RELEASE; ORAL

LAMOTRIGINE

RUBICON	25MG	A202887	001	Jun 17, 2013
	50MG	A202887	002	Jun 17, 2013

LANSOPRAZOLE

CAPSULE, DELAYED REL PELLETS; ORAL

LANSOPRAZOLE

AJANTA PHARMA LTD	15MG	A203957	001	Oct 14, 2016
	30MG	A203957	002	Oct 14, 2016
BRECKENRIDGE	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

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 efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** 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 INC	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			
	SHIRE LLC	EQ 250MG BASE	N021468 001 Oct 26, 2004

LAPYRIUM CHLORIDE; UNDECOYLUM CHLORIDE IODINE COMPLEX

SOLUTION; TOPICAL			
VIRAC REX			
	CHESEBROUGH PONDS	0.5%; 1.8%	N011914 001

LATANOPROST

SOLUTION/DROPS; OPHTHALMIC			
LATANOPROST			
	APOTEX INC	0.005%	A077697 001 Mar 22, 2011

LEFLUNOMIDE

TABLET; ORAL			
LEFLUNOMIDE			
	FOSUN PHARMA	10MG	A077087 001 Sep 13, 2005
		20MG	A077087 002 Sep 13, 2005
	SANDOZ	10MG	A077085 001 Sep 13, 2005
		20MG	A077085 002 Sep 13, 2005

LEPIRUDIN RECOMBINANT

INJECTABLE; INJECTION			
REFLUDAN			
	BAYER HLTHCARE	50MG/VIAL	N020807 001 Mar 06, 1998

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
	FRESENIUS KABI USA	2.5MG	A090491 001 Jun 03, 2011
	IMPAX LABS	2.5MG	A091638 001 Jun 03, 2011
	LANNETT CO INC	2.5MG	A091098 001 Jun 03, 2011
		2.5MG	A202048 001 Oct 29, 2014
	MYLAN	2.5MG	A078190 001 Dec 24, 2008
	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 3MG BASE/ML	A089352 001 Jun 01, 1988
		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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

LEUCOVORIN CALCIUM

INJECTABLE; INJECTION

LEUCOVORIN CALCIUM

+	HOSPIRA	EQ 3MG BASE/ML **	N008107 001	
+		EQ 50MG BASE/VIAL **	N008107 002	
+		EQ 100MG BASE/VIAL **	N008107 004	May 23, 1988
+		EQ 350MG BASE/VIAL **	N008107 005	Apr 05, 1989
	INGENUS PHARMS LLC	EQ 10MG BASE/ML	A210917 001	Nov 23, 2018
	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

LEUCOVORIN CALCIUM PRESERVATIVE FREE

	AM REGENT	EQ 50MG BASE/VIAL	A040338 001	Jan 31, 2001
	HOSPIRA	EQ 10MG BASE/ML **	A040147 001	Jun 25, 1997
	TEVA PARENTERAL	EQ 10MG BASE/ML	A040332 001	Jun 28, 1999

WELLCOVORIN

	GLAXOSMITHKLINE	EQ 5MG BASE/ML	A087439 001	Oct 19, 1982
		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

TABLET; ORAL

LEUCOVORIN CALCIUM

	ANI PHARMS INC	EQ 15MG BASE	A075327 001	Mar 24, 1999
	EPIC PHARMA LLC	EQ 5MG BASE	A074544 001	Aug 28, 1997
		EQ 25MG BASE	A074544 002	Aug 28, 1997
	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
	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

FOR SUSPENSION; INTRAMUSCULAR

LUTRATE DEPOT KIT

+	GP-PHARM SA	22.5MG/VIAL	N205054 001	Aug 28, 2018
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IMPLANT; IMPLANTATION

VIADUR

	ORTHO MCNEIL JANSSEN	EQ 65MG BASE	N021088 001	Mar 03, 2000
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INJECTABLE; INJECTION

LEUPROLIDE ACETATE

	GENZYME	1MG/0.2ML	A075721 001	Nov 29, 2001
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LUPRON

+	ABBVIE ENDOCRINE INC	1MG/0.2ML **	N019010 001	Apr 09, 1985
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LUPRON DEPOT

+	ABBVIE ENDOCRINE INC	3.75MG/VIAL **	N020011 001	Oct 22, 1990
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LUPRON DEPOT-PED

+	ABBVIE ENDOCRINE INC	3.75MG/VIAL, 7.5MG/VIAL **	N020263 003	Apr 16, 1993
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+		7.5MG/VIAL, 7.5MG/VIAL **	N020263 004	Apr 16, 1993
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LEVALBUTEROL HYDROCHLORIDE

SOLUTION; INHALATION

LEVALBUTEROL HYDROCHLORIDE

	MYLAN SPECIALITY LP	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

LEVALLORPHAN TARTRATE

INJECTABLE; INJECTION

LORFAN

	ROCHE	1MG/ML	N010423 001	
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DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

LEVAMISOLE HYDROCHLORIDE

TABLET; ORAL

ERGAMISOL

JANSSEN PHARMA	EQ 50MG BASE	N020035 001	Jun 18, 1990
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LEVETIRACETAM

INJECTABLE; INTRAVENOUS

LEVETIRACETAM

AKORN	500MG/5ML (100MG/ML)	A209934 001	May 04, 2018
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AM REGENT	500MG/5ML (100MG/ML)	A202143 001	Jan 31, 2012
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SOLUTION; ORAL

LEVETIRACETAM

ACI HEALTHCARE LTD	100MG/ML	A078582 001	Jan 15, 2009
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APOTEX INC	100MG/ML	A090187 001	Aug 05, 2011
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TABLET; ORAL

LEVETIRACETAM

ACTAVIS LABS FL INC	250MG	A077408 001	Mar 02, 2009
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	500MG	A077408 002	Mar 02, 2009
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	750MG	A077408 003	Mar 02, 2009
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FOSUN PHARMA	250MG	A077324 001	Jan 15, 2009
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	500MG	A077324 002	Jan 15, 2009
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	750MG	A077324 003	Jan 15, 2009
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	1GM	A077324 004	Jan 15, 2009
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MYLAN	250MG	A078731 001	Feb 10, 2009
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	500MG	A078731 002	Feb 10, 2009
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	750MG	A078731 003	Feb 10, 2009
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	1GM	A078731 004	Feb 10, 2009
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WATSON LABS INC	250MG	A078797 002	Jan 15, 2009
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	500MG	A078797 003	Jan 15, 2009
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	750MG	A078797 004	Jan 15, 2009
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	1GM	A078797 001	Jan 15, 2009
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TABLET, EXTENDED RELEASE; ORAL

LEVETIRACETAM

MYLAN PHARMS INC	500MG	A200475 001	Dec 19, 2011
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	750MG	A200475 002	Dec 19, 2011
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	1GM	A200475 003	Dec 07, 2015
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SANDOZ	500MG	A091668 001	Nov 01, 2012
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	750MG	A091668 002	Nov 01, 2012
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SUN PHARM INDUSTRIES	500MG	A091285 001	Sep 12, 2011
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	750MG	A091285 002	Sep 12, 2011
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VIRTUS PHARMS	500MG	A091291 001	Sep 12, 2011
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	750MG	A091291 002	Sep 12, 2011
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LEVOBETAXOLOL HYDROCHLORIDE

SUSPENSION/DROPS; OPHTHALMIC

BETAXON

ALCON PHARMS LTD	EQ 0.5% BASE	N021114 001	Feb 23, 2000
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LEVOBUNOLOL HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

LEVOBUNOLOL HYDROCHLORIDE

ALCON LABS INC	0.25%	A074851 001	Oct 28, 1996
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APOTEX INC	0.25%	A075473 001	Aug 03, 2000
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	0.5%	A075475 001	Aug 03, 2000
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BAUSCH AND LOMB	0.25%	A074307 001	Mar 04, 1994
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SANDOZ INC	0.5%	A074850 001	Oct 28, 1996
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LEVOBUPIVACAINE HYDROCHLORIDE

INJECTABLE; INJECTION

CHIROCAINE

PURDUE PHARMA LP	EQ 2.5MG BASE/ML	N020997 001	Aug 05, 1999
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	EQ 5MG BASE/ML	N020997 002	Aug 05, 1999
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	EQ 7.5MG BASE/ML	N020997 003	Aug 05, 1999
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LEVOCABASTINE HYDROCHLORIDE

SUSPENSION/DROPS; OPHTHALMIC

LIVOSTIN

NOVARTIS	EQ 0.05% BASE	N020219 001	Nov 10, 1993
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DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

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

LEVOCETIRIZINE DIHYDROCHLORIDE

SOLUTION; ORAL

LEVOCETIRIZINE DIHYDROCHLORIDE

APOTEX 2.5MG/5ML

A202915 001 Aug 21, 2014

TABLET; ORAL

LEVOCETIRIZINE DIHYDROCHLORIDE

APOTEX 5MG

A203027 001 Feb 13, 2015

FOSUN PHARMA 5MG

A090486 001 Mar 26, 2013

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

EMCURE PHARMS LTD EQ 500MG/20ML (EQ 25MG/ML)

A202590 001 Jan 24, 2013

EQ 750MG/30ML (EQ 25MG/ML)

A202590 002 Jan 24, 2013

HOSPIRA INC EQ 500MG/20ML (EQ 25MG/ML)

A078577 001 Aug 12, 2015

EQ 750MG/30ML (EQ 25MG/ML)

A078577 002 Aug 12, 2015

MYLAN ASI 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

SOLUTION; ORAL

LEVAQUIN

+ JANSSEN PHARMS 250MG/10ML

N021721 001 Oct 21, 2004

SOLUTION/DROPS; OPHTHALMIC

IQIIX

+ SANTEN 1.5% **

N021571 001 Mar 01, 2004

LEVOFLOXACIN

APOTEX INC 0.5%

A078282 001 Dec 20, 2010

QUIXIN

+ SANTEN 0.5% **

N021199 001 Aug 18, 2000

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

LEVOFLOXACIN

TABLET; ORAL

LEVAQUIN

+	JANSSEN PHARMS	250MG	N020634	001	Dec 20, 1996
+		500MG	N020634	002	Dec 20, 1996
+		750MG	N020634	003	Sep 08, 2000

LEVOFLOXACIN

MYLAN

		250MG	A076276	001	Jun 20, 2011
		500MG	A076276	002	Jun 20, 2011
		750MG	A077097	001	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 CALCIUM

POWDER; INTRAVENOUS

LEVOLEUCOVORIN CALCIUM

+	ACTAVIS LLC	EQ 175MG BASE/VIAL	N208723	001	Sep 29, 2016
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SOLUTION; INTRAVENOUS

FUSILEV

+	ACROTECH	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

MYLAN TEORANTA

		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

LEVOMEPRMAZINE

INJECTABLE; INJECTION

LEVOPROME

IMMUNEX

		20MG/ML	N015865	001	
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LEVOMETHADYL ACETATE HYDROCHLORIDE

CONCENTRATE; ORAL

ORLAAM

+	ROXANE	10MG/ML **	N020315	001	Jul 09, 1993
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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

LEVONORDEFIN; MEPIVACAINE HYDROCHLORIDE

INJECTABLE; INJECTION

ARESTOCAINE HYDROCHLORIDE W/ LEVONORDEFIN

	SOLVAY	0.05MG/ML; 2%	A085010	001	
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CARBOCAINE W/ NEO-COBEFRIN

	EASTMAN KODAK	0.05MG/ML; 2%	N012125	002	
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ISOCAINE HYDROCHLORIDE W/ LEVONORDEFIN

	SEPTODONT INC	0.05MG/ML; 2%	A084697	001	
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MEPIVACAINE HYDROCHLORIDE W/ LEVONORDEFIN

	BELMORA LLC	0.05MG/ML; 2%	A084850	002	Oct 21, 1983
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POLOCAINE W/ LEVONORDEFIN

	DENTSPLY PHARM	0.05MG/ML; 2%	A089517	001	Apr 14, 1988
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LEVONORDEFIN; PROCAINE HYDROCHLORIDE; PROPOXYCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

RAVOCAINE AND NOVOCAIN W/ NEO-COBEFRIN

	EASTMAN KODAK	0.05MG/ML; 2%; 0.4%	N008592	007	
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LEVONORGESTREL

IMPLANT; IMPLANTATION

JADELLE

+	POPULATION COUNCIL	75MG/IMPLANT **	N020544	001	Nov 01, 1996
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LEVONORGESTREL

WYETH PHARMS INC

		75MG/IMPLANT	N020627	001	Aug 15, 1996
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NORPLANT

POPULATION COUNCIL

		36MG/IMPLANT	N019897	001	Dec 10, 1990
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Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

LEVONORGESTREL

IMPLANT; IMPLANTATION

NORPLANT SYSTEM IN PLASTIC CONTAINER

WYETH PHARMS INC 36MG/IMPLANT

N020088 001 Dec 10, 1990

TABLET; ORAL

LEVONORGESTREL

ACCORD HLTHCARE 1.5MG

A207660 001 May 02, 2019

ALVOGEN 1.5MG

A202246 001 Jun 05, 2015

FDN CONSUMER 0.75MG **

A078665 001 Aug 28, 2009

1.5MG

A200670 001 Jul 12, 2012

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

PLAN B

+ FDN CONSUMER 0.75MG **

N021045 001 Jul 28, 1999

+ 0.75MG **

N021045 002 Aug 24, 2006

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

SENTYNL THERAPS INC 1MG

A074278 002 Jun 18, 2018

LEVOTHYROXINE SODIUM

POWDER; INTRAVENOUS

LEVOTHYROXINE SODIUM

PAR STERILE PRODUCTS 200MCG/VIAL

A205366 001 Dec 07, 2015

TABLET; ORAL

EUTHYROX

PROVELL 0.3MG

N021292 012 May 31, 2002

LEVOLET

GENUS LIFESCIENCES 0.025MG

N021137 001 Jun 06, 2003

0.05MG

N021137 002 Jun 06, 2003

0.075MG

N021137 003 Jun 06, 2003

0.088MG

N021137 004 Jun 06, 2003

0.1MG

N021137 005 Jun 06, 2003

0.112MG

N021137 006 Jun 06, 2003

0.125MG

N021137 007 Jun 06, 2003

0.137MG

N021137 008 Jun 06, 2003

0.15MG

N021137 009 Jun 06, 2003

0.175MG

N021137 010 Jun 06, 2003

0.2MG

N021137 011 Jun 06, 2003

0.3MG

N021137 012 Jun 06, 2003

LEVOTHYROXINE SODIUM

AMNEAL PHARMS LLC 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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

6-256(of 430)

** See List Footnote

LEVOTHYROXINE SODIUM

TABLET; ORAL

LEVOTHYROXINE SODIUM

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

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

XYLOCAINE

+ ASTRAZENECA

5% **

N008048 001

PATCH; TOPICAL

DENTIPATCH

NOVEN

46.1MG/PATCH

N020575 002 May 21, 1996

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

AKORN

1%

A085037 001

2%

A085037 002

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

GD SEARLE LLC

1%

A083135 001

2%

A083135 002

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

6-257(of 430)

** See List Footnote

LIDOCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

LIDOCAINE HYDROCHLORIDE

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	
	2GM/VIAL	N018543 002	
LUITPOLD	2%	A083198 001	
LYPHOMED	1%	A080390 001	
	2%	A080390 002	
MILES	1%	A080414 001	
	2%	A080414 002	
MYLAN LABS LTD	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	
WEST-WARD PHARMS INT	1%	A080407 001	
	2%	A080407 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			
INTL MEDICATION	4%	N017702 002	
	20%	N017702 001	
MYLAN LABS LTD	2%	A090665 001	Sep 27, 2010
WEST-WARD PHARMS INT	1%	A084625 001	
	2%	A084625 002	
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	

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

6-258(of 430)

** See List Footnote

LIDOCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

XYLOCAINE DENTAL

DENTSPLY PHARM 2% N021380 001

XYLOCAINE PRESERVATIVE FREE

+ FRESENIUS KABI USA 1% N016801 005 Jan 19, 1988

+ 2% N016801 001

+ 4% N016801 002

+ 10% N016801 003

+ 20% N016801 004

INJECTABLE; SPINAL

XYLOCAINE 1.5% W/ DEXTROSE 7.5%

FRESENIUS KABI USA 1.5% N016297 001

XYLOCAINE 5% W/ GLUCOSE 7.5%

ASTRAZENECA 5% N010496 002 Jul 07, 1982

JELLY; TOPICAL

ANESTACON

BIONPHARMA INC 2% A080429 001

LIDOCAINE HYDROCHLORIDE

ACP NIMBLE 2% A081318 001 Apr 29, 1993

WATSON LABS INC 2% A040837 001 Mar 23, 2011

SOLUTION; ORAL

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

WOCKHARDT BIO AG 4% A087881 001 Nov 18, 1982

LTA II KIT

HOSPIRA 4% A080409 001

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

DISC; TOPICAL

EMLA

ASTRAZENECA 2.5%; 2.5% N020962 001 Feb 04, 1998

LINCOMYCIN HYDROCHLORIDE

CAPSULE; ORAL

LINCOCIN

PHARMACIA AND UPJOHN EQ 250MG BASE N050316 001

EQ 500MG BASE N050316 002

INJECTABLE; INJECTION

LINCOMYCIN HYDROCHLORIDE

WATSON LABS 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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

6-259(of 430)

** See List Footnote

LINDANE

LOTION; TOPICAL

KWELL

REED AND CARNRICK

1%

A084218 002

1%

N006309 003

LINDANE

WOCKHARDT

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

SCABENE

STIEFEL

1%

A087940 001 Apr 08, 1983

LINEZOLID

SOLUTION; INTRAVENOUS

ZYVOX

+ PHARMACIA AND UPJOHN

400MG/200ML (2MG/ML) **

N021131 002 Apr 18, 2000

TABLET; ORAL

LINEZOLID

MYLAN

600MG

A078845 001 Dec 21, 2015

ZYVOX

+ PHARMACIA AND UPJOHN

400MG **

N021130 001 Apr 18, 2000

LIOTHYRONINE SODIUM

TABLET; ORAL

LIOTHYRONINE SODIUM

MYLAN

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

TEVA PHARMS USA

EQ 0.005MG BASE

A211510 001 Oct 26, 2018

EQ 0.025MG BASE

A211510 002 Oct 26, 2018

EQ 0.05MG BASE

A211510 003 Oct 26, 2018

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

LISINAPRIL

TABLET; ORAL

LISINAPRIL

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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

LISINAPRILTABLET; ORAL
LISINAPRIL

	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
INVATECH	2.5MG	A075903 001	Jul 01, 2002
	2.5MG	A075999 001	Jul 01, 2002
	5MG	A075903 002	Jul 01, 2002
	5MG	A075999 002	Jul 01, 2002
	10MG	A075903 003	Jul 01, 2002
	10MG	A075999 003	Jul 01, 2002
	20MG	A075903 004	Jul 01, 2002
	20MG	A075999 004	Jul 01, 2002
	30MG	A075903 005	Jul 01, 2002
	30MG	A075999 005	Jul 01, 2002
	40MG	A075903 006	Jul 01, 2002
	40MG	A075999 006	Jul 01, 2002
MYLAN	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
	10MG	N019558 002	Dec 29, 1987
	20MG	N019558 003	Dec 29, 1987

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
MYLAN	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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

6-261(of 430)

** See List Footnote

LITHIUM CARBONATE

TABLET, EXTENDED RELEASE;ORAL

LITHIUM CARBONATE

ABLE	300MG	A076382 001	Apr 21, 2003
HERITAGE PHARMA	300MG	A076170 001	Jun 10, 2002
	450MG	A076366 001	Aug 21, 2003
HIKMA INTL PHARMS	450MG	A076490 001	Jun 17, 2003

LITHIUM CITRATE

SYRUP;ORAL

LITHONATE

SOLVAY	EQ 300MG CARBONATE/5ML	N017672 001	
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LOMEFLOXACIN HYDROCHLORIDE

TABLET;ORAL

MAXAQUIN

PHARMACIA	EQ 400MG BASE	N020013 001	Feb 21, 1992
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LOMUSTINE

CAPSULE;ORAL

GLEOSTINE

+ CORDEN PHARMA	5MG	N017588 004	Dec 19, 2014
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LOPERAMIDE HYDROCHLORIDE

CAPSULE;ORAL

IMODIUM

J AND J CONSUMER INC	2MG **	N017690 001	
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+ J AND J CONSUMER INC	2MG **	N017694 001	
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LOPERAMIDE HYDROCHLORIDE

ROXANE	2MG	A073080 001	Nov 27, 1991
TEVA	2MG	A073122 001	Aug 30, 1991
YAOPHARMA CO LTD	2MG	A072993 001	Aug 28, 1992

SOLUTION;ORAL

IMODIUM

JANSSEN PHARMS	1MG/5ML	N019037 001	Jul 31, 1984
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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
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
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LOPERAMIDE HYDROCHLORIDE; SIMETHICONE

TABLET, CHEWABLE;ORAL

IMODIUM MULTI-SYMPOM RELIEF

+ J AND J CONSUMER INC	2MG;125MG	N020606 001	Jun 26, 1996
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LOPINAVIR; RITONAVIR

CAPSULE;ORAL

KALETRA

ABBVIE	133.3MG;33.3MG	N021226 001	Sep 15, 2000
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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

DISCONTINUED DRUG PRODUCT LIST

6-262(of 430)

** See List Footnote

LORATADINE

CAPSULE; ORAL

LORATADINE

BIONPHARMA INC 10MG A202538 001 Dec 21, 2018

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

TABLET; ORAL

LORATADINE

MYLAN 10MG A075790 001 Nov 07, 2008

10MG A078447 001 Aug 12, 2011

PERRIGO 10MG N021512 001 Jun 24, 2004

LORATADINE; PSEUDOEPHEDRINE SULFATE

TABLET, EXTENDED RELEASE; ORAL

LORATADINE AND PSEUDOEPHEDRINE SULFATE

HERITAGE PHARMA 5MG;120MG A076208 001 Jan 28, 2004

LORAZEPAM

INJECTABLE; INJECTION

LORAZEPAM

AKORN 2MG/ML A074974 001 Jul 23, 1998

BEDFORD 2MG/ML A077076 001 Jul 13, 2005

4MG/ML A077076 002 Jul 13, 2005

DAVA PHARMS INC 2MG/ML A074793 001 Mar 16, 2000

4MG/ML A074793 002 Mar 16, 2000

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

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

MYLAN ASI 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

WEST-WARD PHARMS INT 2MG/ML A074496 001 Sep 28, 1998

4MG/ML A074496 002 Sep 28, 1998

LORAZEPAM PRESERVATIVE FREE

BEDFORD LABS 2MG/ML A077074 001 Jul 13, 2005

4MG/ML A077074 002 Jul 13, 2005

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

ANDA REPOSITORY 0.5MG A072555 002 Mar 29, 1991

1MG A072555 003 Mar 29, 1991

2MG A072555 001 Mar 29, 1991

ANI PHARMS INC 0.5MG A077396 001 Dec 13, 2006

1MG A077396 002 Dec 13, 2006

2MG A077396 003 Dec 13, 2006

HALSEY 0.5MG A071434 001 Sep 01, 1987

1MG A071435 001 Sep 01, 1987

2MG A071436 001 Sep 01, 1987

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

LORAZEPAM

TABLET; ORAL

LORAZEPAM

MUTUAL PHARM	0.5MG	A070472 001	Dec 10, 1985
	1MG	A070473 001	Dec 10, 1985
	2MG	A070474 001	Dec 10, 1985
MYLAN	0.5MG	A071591 002	Oct 13, 1987
	1MG	A071591 003	Oct 13, 1987
	2MG	A071591 001	Oct 13, 1987
PAR PHARM	0.5MG	A070675 001	Dec 01, 1986
	1MG	A070676 001	Dec 01, 1986
	2MG	A070677 001	Dec 01, 1986
SANDOZ	0.5MG	A071193 001	Apr 15, 1988
	1MG	A071194 001	Apr 15, 1988
	2MG	A071195 001	Apr 15, 1988
SUN PHARM INDS LTD	0.5MG	A076045 001	Aug 29, 2001
	1MG	A076045 002	Aug 29, 2001
	2MG	A076045 003	Aug 29, 2001
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

LORCASERIN HYDROCHLORIDE

TABLET; ORAL

BELVIQ

+ EISAI INC	10MG	N022529 001	Jun 27, 2012
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TABLET, EXTENDED RELEASE; ORAL

BELVIQ XR

+ EISAI INC	20MG	N208524 001	Jul 15, 2016
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LOSARTAN POTASSIUM

TABLET; ORAL

COZAAR

+ MERCK SHARP DOHME	25MG	N020386 001	Apr 14, 1995
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+	50MG	N020386 002	Apr 14, 1995
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LOSARTAN POTASSIUM

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
TEVA	25MG	A076958 001	Apr 06, 2010
	50MG	A076958 002	Apr 06, 2010
	100MG	A076958 003	Apr 06, 2010

LOTEPREDNOL ETABONATE

SUSPENSION/DROPS; OPHTHALMIC

LOTEMAX

PHARMOS	0.5%	N020841 001	Mar 09, 1998
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LOVASTATIN

TABLET; ORAL

LOVASTATIN

MYLAN	10MG	A075451 001	Dec 17, 2001
	10MG	A075935 001	Dec 17, 2001
	20MG	A075451 002	Dec 17, 2001
	20MG	A075935 002	Dec 17, 2001

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

LOVASTATINTABLET; ORAL
LOVASTATIN

	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 PHARMA BV	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

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	

LUCINACTANT

SUSPENSION; INTRATRACHEAL

SURFAXIN			
WINDTREE THERAP	8.5ML	N021746 001	Mar 06, 2012

LURASIDONE HYDROCHLORIDE

TABLET; ORAL

LURASIDONE HYDROCHLORIDE			
AMNEAL PHARMS CO	20MG	A208002 001	Jan 03, 2019
	40MG	A208002 002	Jan 03, 2019
	60MG	A208002 003	Jan 03, 2019
	80MG	A208002 004	Jan 03, 2019
	120MG	A208002 005	Jan 03, 2019
EMCURE PHARMS LTD	20MG	A208058 001	Sep 04, 2019
	40MG	A208058 002	Sep 04, 2019
	60MG	A208058 003	Sep 04, 2019
	80MG	A208058 004	Sep 04, 2019
INVAGEN PHARMS	20MG	A208028 001	Jan 03, 2019
	40MG	A208028 002	Jan 03, 2019
	60MG	A208028 003	Jan 03, 2019
	80MG	A208028 004	Jan 03, 2019
	120MG	A208028 005	Jan 03, 2019
LUPIN LTD	20MG	A208031 001	Jan 03, 2019
	40MG	A208031 002	Jan 03, 2019
	60MG	A208031 003	Jan 03, 2019
	80MG	A208031 004	Jan 03, 2019
	120MG	A208031 005	Jan 03, 2019
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
TORRENT	20MG	A208055 001	Jan 03, 2019
	40MG	A208055 002	Jan 03, 2019

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

LURASIDONE HYDROCHLORIDE

TABLET; ORAL

LURASIDONE HYDROCHLORIDE

80MG	A208055 003	Jan 03, 2019
120MG	A208055 004	Jan 03, 2019
ZYDUS PHARMS	20MG	A208052 001
	40MG	A208052 002
	60MG	A208052 003
	80MG	A208052 004
	120MG	A208052 005

LUTROPIN ALFA

INJECTABLE; SUBCUTANEOUS

LUVERIS

EMD SERONO	75 IU/VIAL	N021322 001	Oct 08, 2004
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LYPRESSIN

SOLUTION; NASAL

DIAPID

NOVARTIS	0.185MG/ML	N016755 001
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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
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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
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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
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SOLUTION; IRRIGATION

PHYSIOSOL IN PLASTIC CONTAINER

HOSPIRA INC	14MG/100ML; 37MG/100ML; 222MG/100ML; 526MG/100ML; 502MG/100ML	N018406 001
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PHYSIOSOL PH 7.4 IN PLASTIC CONTAINER

HOSPIRA INC	30MG/100ML; 37MG/100ML; 222MG/100ML; 526MG/100ML; 502MG/100ML	N018406 002	Jul 08, 1982
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SYNOVALYTE IN PLASTIC CONTAINER

BAXTER HLTHCARE	30MG/100ML; 37MG/100ML; 368MG/100ML; 526MG/100ML; 502MG/100ML	N019326 001	Jan 25, 1985
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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; 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, 2.86GM, 5.6GM **	N203595 001	Jan 18, 2013
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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
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Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

MALATHION

LOTION; TOPICAL

MALATHION

MYLAN PHARMS INC 0.5%

A078743 001 Mar 06, 2009

OVIDE

+ TARO PHARM INDS LTD 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

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

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

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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

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
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
WATSON LABS TEVA	50MG	A072163 001	Jun 01, 1988

MASOPROCOL

CREAM; TOPICAL

ACTINEX

UNIV AZ CANCER CTR	10%	N019940 001	Sep 04, 1992
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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	
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MECAMYLAMINE HYDROCHLORIDE

TABLET; ORAL

INVERSINE

+ TARGACEPT	2.5MG **	N010251 001	
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MECASERMIN RINFABATE RECOMBINANT

INJECTABLE; SUBCUTANEOUS

IPLEX

INSMED	36MG/0.6ML	N021884 001	Dec 12, 2005
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MECHLORETHAMINE HYDROCHLORIDE

INJECTABLE; INJECTION

MUSTARGEN

+ RECORDATI RARE	10MG/VIAL	N006695 001	
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MECLIZINE HYDROCHLORIDE

TABLET; ORAL

MECLIZINE HYDROCHLORIDE

ABC HOLDING	12.5MG	A085253 001	
	25MG	A085252 001	
AMNEAL PHARMS	50MG	A201451 003	Feb 23, 2011
ANABOLIC	25MG	A085891 001	
ANI PHARMS INC	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	
IVAX SUB TEVA PHARMS	12.5MG	A083784 001	
KV PHARM	12.5MG	A085524 001	

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

MECLIZINE HYDROCHLORIDE

TABLET; ORAL

MECLIZINE HYDROCHLORIDE

	25MG	A085523	001	
MYLAN PHARMS INC	12.5MG	A202640	001	Sep 17, 2012
	25MG	A202640	002	Sep 17, 2012
	50MG	A202640	003	Sep 17, 2012
PAR PHARM	50MG	A089674	001	Mar 31, 1988
PLIVA	25MG	A088734	001	Dec 11, 1985
RISING	12.5MG	A040179	001	Jan 30, 1997
	25MG	A040179	002	Jan 30, 1997
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

IVAX SUB TEVA PHARMS	25MG	A084976	001	
NEXGEN PHARMA INC	25MG	A086392	001	
PLIVA	25MG	A088733	001	Dec 11, 1985

MECLOCYCLINE SULFOSALICYLATE

CREAM; TOPICAL

MECLAN

JOHNSON AND JOHNSON	1%	N050518	001	
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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 INC	EQ 50MG BASE	A071468	001	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
FOSUN PHARMA	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

+ PHARMACIA AND UPJOHN	100MG/ML **	N012541	002	
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MEDROXYPROGESTERONE ACETATE

CIPLA	150MG/ML	A210335	001	Jan 25, 2019
SANDOZ INC	150MG/ML	A078711	001	May 20, 2009
TEVA PHARMS USA	150MG/ML	A076552	001	Oct 27, 2004

TABLET; ORAL

AMEN

AMARIN PHARMS	10MG	A083242	001	
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CURRETAB

SOLVAY	10MG	A085686	001	
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Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

MEDROXYPROGESTERONE ACETATE

TABLET; ORAL

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

MEDRYSONE

SUSPENSION; OPHTHALMIC

HMS

ALLERGAN	1%	N016624	003	
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MEFENAMIC ACID

CAPSULE; ORAL

MEFENAMIC ACID

BRECKENRIDGE	250MG	A090359	001	Feb 05, 2013
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MEFLOQUINE HYDROCHLORIDE

TABLET; ORAL

LARIAM

+ ROCHE	250MG **	N019591	001	May 02, 1989
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MEFLOQUINE HYDROCHLORIDE

HIKMA	250MG	A076523	001	Oct 01, 2004
HIKMA INTL PHARMS	250MG	A077699	001	Apr 21, 2010
SANDOZ	250MG	A076175	001	Feb 20, 2002
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
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MEGESTROL ACETATE

HIKMA	40MG/ML	A075997	001	Feb 15, 2002
PHARM ASSOC	40MG/ML	A077404	001	Feb 16, 2006

TABLET; ORAL

MEGACE

+ BRISTOL MYERS SQUIBB	20MG **	N016979	001	
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+ BRISTOL MYERS SQUIBB	40MG **	N016979	002	
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MEGESTROL ACETATE

TEVA	40MG	A074745	001	Feb 27, 1998
USL PHARMA	20MG	A070646	001	Oct 02, 1987
	40MG	A070647	001	Oct 02, 1987

MELOXICAM

SUSPENSION; ORAL

MOBIC

+ AVONDALE PHARMS	7.5MG/5ML **	N021530	001	Jun 01, 2004
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TABLET; ORAL

MELOXICAM

ANDA REPOSITORY	7.5MG	A077935	001	Jul 19, 2006
	15MG	A077935	002	Jul 19, 2006
CR DOUBLE CRANE	7.5MG	A078039	001	Dec 14, 2006
	15MG	A078039	002	Dec 14, 2006
HERITAGE PHARMA	7.5MG	A077936	001	Jul 19, 2006
	15MG	A077936	002	Jul 19, 2006
IMPAX LABS INC	7.5MG	A077930	001	Jul 19, 2006
	15MG	A077930	002	Jul 19, 2006
MYLAN	7.5MG	A077923	001	Jul 19, 2006
	7.5MG	A077934	001	Jul 20, 2006
	15MG	A077923	002	Jul 19, 2006
	15MG	A077934	002	Jul 20, 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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

MELPHALAN HYDROCHLORIDE

INJECTABLE; INJECTION

MELPHALAN HYDROCHLORIDE

MYLAN INSTITUTIONAL	EQ 50MG BASE/VIAL	A090299 001	Oct 27, 2009
PAR STERILE PRODUCTS	EQ 50MG BASE/VIAL	A204773 001	Aug 22, 2016
TWI PHARMS	EQ 50MG BASE/VIAL	A211463 001	Sep 13, 2019
US WORLDMEDS LLC	EQ 50MG BASE/VIAL	A207032 001	May 03, 2019

MEMANTINE HYDROCHLORIDE

SOLUTION; ORAL

MEMANTINE HYDROCHLORIDE

TORRENT	2MG/ML	A205446 001	Dec 07, 2015
NAMENDA + ALLERGAN	2MG/ML	N021627 001	Apr 18, 2005

TABLET; ORAL

MEMANTINE HYDROCHLORIDE

APOTEX INC	5MG	A090244 001	Jul 11, 2018
	10MG	A090244 002	Jul 11, 2018
MYLAN	5MG	A079225 001	Jan 30, 2015
	10MG	A079225 002	Jan 30, 2015
ORCHID HLTHCARE	5MG	A090044 001	Mar 12, 2012
	10MG	A090044 002	Mar 12, 2012

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	
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MENADIONE

TABLET; ORAL

MENADIONE

LILLY	5MG	N002139 003	
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MENOTROPINS (FSH; LH)

INJECTABLE; INJECTION

HUMEGON

ORGANON USA INC	75 IU/VIAL; 75 IU/VIAL	N020328 001	Sep 01, 1994
	150 IU/VIAL; 150 IU/VIAL	N020328 002	Sep 01, 1994

MENOTROPINS

FERRING	75 IU/VIAL; 75 IU/VIAL	A073598 001	Jan 30, 1997
	150 IU/VIAL; 150 IU/VIAL	A073599 001	Jan 30, 1997

PERGONAL

SERONO	75 IU/AMP; 75 IU/AMP	N017646 001	
	150 IU/AMP; 150 IU/AMP	N017646 002	May 20, 1985

REPRONEX

FERRING	150 IU/VIAL; 150 IU/VIAL	N021047 002	Aug 27, 1999
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INJECTABLE; INTRAMUSCULAR, SUBCUTANEOUS

REPRONEX

FERRING	75 IU/VIAL; 75 IU/VIAL	N021047 001	Aug 27, 1999
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MEPENZOLATE BROMIDE

SOLUTION; ORAL

CANTIL

SANOFI AVENTIS US	25MG/5ML	N010679 004	
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TABLET; ORAL

CANTIL

+ SANOFI AVENTIS US	25MG	N010679 003	
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MEPERIDINE HYDROCHLORIDE

INJECTABLE; INJECTION

DEMEROL

US PHARM HOLDINGS	25MG/ML	N005010 007	
	50MG/ML	N005010 002	
	75MG/ML	N005010 009	
	100MG/ML	N005010 003	

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

MEPERIDINE HYDROCHLORIDE

INJECTABLE; INJECTION

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	10MG/ML	A086332	001	
PARKE DAVIS	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
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				
US PHARM HOLDINGS	50MG/5ML **	N005010	005	
TABLET; ORAL				
DEMEROL				
+ US PHARM HOLDINGS	50MG	N005010	001	
+	100MG	N005010	004	
MEPERIDINE HYDROCHLORIDE				
ANDA REPOSITORY	50MG	A040893	001	Jun 24, 2009
	75MG	A040893	002	Jun 24, 2009
	100MG	A040893	003	Jun 24, 2009
	150MG	A040893	004	Jun 24, 2009
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
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
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
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DISCONTINUED DRUG PRODUCT LIST

6-272(of 430)

** See List Footnote

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
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MEPIVACAINE HYDROCHLORIDE

INJECTABLE; INJECTION

ARESTOCAINE HYDROCHLORIDE

SOLVAY	3%	A084777	002	Apr 18, 1982
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CARBOCAINE

+ EASTMAN KODAK	3% **	N012125	003
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ISOCAINE HYDROCHLORIDE

SEPTODONT INC	3%	A080925	001
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MEPIVACAINE HYDROCHLORIDE

BELMORA LLC	3%	A083559	001
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HOSPIRA INC	3%	A040806	001	Apr 28, 2008
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INTL MEDICATION SYS	1%	A087509	001	Oct 05, 1982
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WATSON LABS	1%	A088769	001	Nov 20, 1984
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	2%	A088770	001	Nov 20, 1984
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POLOCAINE

DENTSPLY PHARM	3%	A088653	001	Aug 21, 1984
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MEPREDNISONE

TABLET; ORAL

BETAPAR

SCHERING	4MG	N016053	002
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MEPROBAMATE

CAPSULE; ORAL

EQUANIL

WYETH AYERST	400MG	N012455	002
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CAPSULE, EXTENDED RELEASE; ORAL

MEPROSPAN

MEDPOINTE PHARM HLC	200MG	N011284	001
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	400MG	N011284	002
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TABLET; ORAL

AMOSENE

FERNDAL LABS	400MG	A084030	001
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BAMATE

ALRA	200MG	A080380	001
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	400MG	A080380	002
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EQUANIL

WYETH AYERST	200MG	N010028	005
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	400MG	N010028	004
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MEPRIAM

TEVA	400MG	N016069	001
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MEPROBAMATE

AUROLIFE PHARMA LLC	400MG	A080655	001
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BARR	600MG	A084230	001
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ELKINS SINN	200MG	N015426	002
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	400MG	N015426	001
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HEATHER	400MG	N016928	003
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	600MG	A084329	001
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IMPAX LABS	200MG	N014322	002
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	400MG	N014322	001
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IVAX SUB TEVA PHARMS	200MG	N015438	001
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	400MG	N015438	002
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	600MG	A084181	001
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IVC INDS	400MG	A084153	001
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LANNETT	200MG	N014882	002
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	400MG	N014882	001
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LEDERLE	400MG	A086299	001
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LEE KM	400MG	A089538	001	Nov 25, 1987
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MALLARD	400MG	N015072	002
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MK LABS	200MG	N014368	004
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Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

MEPROBAMATE

TABLET; ORAL

MEPROBAMATE

	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	
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	
	400MG		N015139	005	
VANGARD	400MG		A088011	001	Jul 14, 1982
WATSON LABS	200MG		A085720	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
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MEROPENEM

INJECTABLE; INJECTION

MEROPENEM

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	
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DISCONTINUED DRUG PRODUCT LIST

6-274(of 430)

** See List Footnote

MESALAMINE

ENEMA;RECTAL				
MESALAMINE				
G AND W LABS INC	4GM/60ML		A076841 001	Sep 30, 2004
SUPPOSITORY;RECTAL				
CANASA				
ALLERGAN	500MG		N021252 001	Jan 05, 2001
ROWASA				
+ MEDA PHARMS	500MG **		N019919 001	Dec 18, 1990
TABLET, DELAYED RELEASE;ORAL				
ASACOL				
APIL	400MG		N019651 001	Jan 31, 1992

MESNA

INJECTABLE;INTRAVENOUS				
MESNA				
MYLAN INSTITUTIONAL	100MG/ML		A076488 001	Mar 08, 2012
MYLAN LABS LTD	100MG/ML		A203364 001	Jul 18, 2014

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	

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

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
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ENOVID-E

GD SEARLE LLC	0.1MG; 2.5MG	N010976 006
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TABLET; ORAL-21

ENOVID-E 21

GD SEARLE LLC	0.1MG; 2.5MG	N010976 007
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METAPROTERENOL SULFATE

AEROSOL, METERED; INHALATION

ALUPENT

BOEHRINGER INGELHEIM	0.65MG/INH	N016402 001
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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
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DEY

0.6%	A071018 001	Jul 27, 1988
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0.33%	A071806 001	Aug 05, 1988
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0.5%	A071805 001	Aug 05, 1988
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5%	A070805 001	Aug 17, 1987
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MYLAN SPECIALITY LP

0.4%	A071786 001	Aug 05, 1988
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0.6%	A070804 001	Aug 17, 1987
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NEPHRON

0.4%	A071855 001	Jul 14, 1988
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0.6%	A071726 001	Jul 14, 1988
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WOCKHARDT

0.4%	A075586 001	May 30, 2002
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0.6%	A075586 002	May 30, 2002
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5%	A072190 001	Jun 07, 1988
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PROMETA

MURO

5%	A073340 001	Mar 30, 1992
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SYRUP; ORAL

ALUPENT

BOEHRINGER INGELHEIM	10MG/5ML	N017571 001
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METAPROTERENOL SULFATE

ACP NIMBLE

10MG/5ML	A072761 001	Feb 27, 1992
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APOTEX INC

10MG/5ML	A075235 001	Jan 27, 2000
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G AND W LABS INC

10MG/5ML	A073034 001	Aug 30, 1991
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MORTON GROVE

10MG/5ML	A071656 001	Oct 13, 1987
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WOCKHARDT

10MG/5ML	A074702 001	Mar 24, 1997
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PROMETA

MURO

10MG/5ML	A072023 001	Sep 15, 1988
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TABLET; ORAL

ALUPENT

BOEHRINGER INGELHEIM	10MG	N015874 002
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	20MG	N015874 001
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METAPROTERENOL SULFATE

AM THERAP

10MG	A072054 001	Jun 23, 1988
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20MG	A072055 001	Jun 23, 1988
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HERITAGE PHARMA

10MG	A072519 001	Mar 30, 1990
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20MG	A072520 001	Mar 30, 1990
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PAR PHARM

10MG	A072024 001	Jun 28, 1988
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20MG	A072025 001	Jun 28, 1988
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USL PHARMA

10MG	A071013 001	Jan 25, 1988
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20MG	A071014 001	Jan 25, 1988
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WATSON LABS

10MG	A073013 001	Jan 31, 1991
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20MG	A072795 001	Jan 31, 1991
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DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

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

METAXALONE

TABLET; ORAL

METAXALONE

+ PRIMUS PHARMS 640MG ** N022503 001 Jun 01, 2015

SKELAXIN

+ KING PHARMS 400MG ** N013217 001

METFORMIN HYDROCHLORIDE

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

BARR 500MG A075971 001 Jan 25, 2002

850MG A075971 002 Jan 25, 2002

1GM A075971 003 Jan 25, 2002

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

MYLAN 500MG A075976 001 Jan 24, 2002

850MG A075976 002 Jan 24, 2002

1GM A075976 003 Jan 24, 2002

MYLAN PHARMS INC 500MG A075969 001 Jan 29, 2002

850MG A075969 002 Jan 29, 2002

1GM A075969 003 Jan 29, 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 INDUSTRIES 500MG A076038 001 Feb 21, 2002

850MG A076038 002 Feb 21, 2002

1GM A076038 003 Feb 21, 2002

SUNSHINE LAKE 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

850MG A076328 002 Dec 16, 2002

1GM A076328 003 Dec 16, 2002

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

+ 1GM ** N021574 002 Apr 27, 2004

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

6-277(of 430)

** See List Footnote

METFORMIN HYDROCHLORIDETABLET, EXTENDED RELEASE;ORAL
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
	BARR	500MG	A076496 001	Nov 25, 2005
	IMPAX LABS	500MG	A076249 001	Jul 30, 2004
		750MG	A076985 001	Sep 13, 2005
	IVAX SUB TEVA PHARMS	500MG	A076545 001	Dec 01, 2003
	MYLAN	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 PHARMS LTD	750MG	A079226 001	Feb 18, 2010
	WATSON LABS INC	500MG	A076818 001	Dec 14, 2004
	YICHANG HUMANWELL	500MG	A211052 001	Sep 24, 2018
		750MG	A211052 002	Sep 24, 2018

METFORMIN HYDROCHLORIDE; PIOGLITAZONE HYDROCHLORIDE

TABLET;ORAL

PIOGLITAZONE HYDROCHLORIDE AND METFORMIN HYDROCHLORIDE

	MYLAN	500MG;EQ 15MG BASE	A090406 001	Feb 25, 2011
		850MG;EQ 15MG BASE	A090406 002	Feb 25, 2011

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

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

METHACHOLINE CHLORIDE

FOR SOLUTION;INHALATION

PROVOCHOLINE

+	METHAPHARM	1600MG/VIAL	N019193 002	Aug 29, 2016
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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
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DISCONTINUED DRUG PRODUCT LIST

6-278(of 430)

** See List Footnote

METHADONE HYDROCHLORIDE

POWDER; FOR RX COMPOUNDING

METHADONE HYDROCHLORIDE

MALLINCKRODT INC

50GM/BOT

N006383 002

100GM/BOT

N006383 003

500GM/BOT

N006383 004

SYRUP; ORAL

DOLOPHINE HYDROCHLORIDE

HIKMA

10MG/30ML

N006134 004

TABLET; ORAL

METHADONE HYDROCHLORIDE

ROXANE

5MG

A088108 001 Mar 08, 1983

10MG

A088109 001 Mar 08, 1983

40MG

A074081 001 Apr 28, 1995

VISTAPHARM

5MG

A040241 001 May 29, 1998

TABLET, DISPERSIBLE; ORAL

WESTADONE

SANDOZ

2.5MG

N017108 001

TABLET, EFFERVESCENT; ORAL

WESTADONE

SANDOZ

5MG

N017108 002

10MG

N017108 003

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

RECORDATI RARE

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

INVATECH

25MG

A040102 001 Aug 28, 1996

50MG

A040102 002 Aug 28, 1996

MICRO LABS

25MG

A207438 001 Oct 05, 2018

50MG

A207438 002 Oct 05, 2018

NEPTAZANE

+ LEDERLE

25MG **

N011721 002 Nov 25, 1991

+

50MG **

N011721 001

METHDILAZINE

TABLET, CHEWABLE; ORAL

TACARYL

WESTWOOD SQUIBB

3.6MG

N011950 009

DISCONTINUED DRUG PRODUCT LIST

6-279(of 430)

** See List Footnote

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

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

METHIMAZOLE

TABLET; ORAL

METHIMAZOLE

ECI PHARMS LLC 15MG A040619 003 Jul 12, 2005

20MG A040547 004 Feb 18, 2005

MYLAN 20MG A040350 003 Jun 07, 2001

SUN PHARM INDS INC 5MG A040870 001 Sep 25, 2007

10MG A040870 002 Sep 25, 2007

TAPAZOLE

+ KING PHARMS 5MG ** N007517 002

+ 10MG ** N007517 004

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

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

AM THERAP 500MG A089417 001 Feb 11, 1987

750MG A089418 001 Feb 11, 1987

ANI PHARMS INC 500MG A084277 001

750MG A084276 002

ASCOT 500MG A087660 001 Oct 27, 1982

750MG A087661 001 Oct 27, 1982

CLONMEL HLTHCARE 500MG A085961 001

750MG A085963 001

FOSUN PHARMA 500MG A084616 001

750MG A084615 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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

METHOCARBAMOL

TABLET; ORAL

METHOCARBAMOL

LANNETT CO INC	500MG	A084756 002	Mar 31, 2003
	750MG	A084756 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
	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			
+ AUXILIUM PHARMS LLC	500MG	N011011 004	

METHOHEXITAL SODIUM

INJECTABLE; INJECTION

BREVITAL SODIUM

PAR STERILE PRODUCTS	200MG/VIAL	N011559 004	Dec 21, 2012
	5GM/VIAL	N011559 003	

METHOTREXATE

SOLUTION; SUBCUTANEOUS

OTREXUP

+ ANTARES PHARMA INC	7.5MG/0.4ML (7.5MG/0.4ML)	N204824 005	Nov 07, 2014
OTREXUP PFS			
+ ANTARES PHARMA INC	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

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
FOLEX PFS			
PHARMACIA AND UPJOHN	EQ 25MG BASE/ML	A081242 001	Aug 23, 1991
	EQ 25MG BASE/ML	A089180 001	Jan 03, 1986

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

METHOTREXATE SODIUM

INJECTABLE; INJECTION

METHOTREXATE LPF

HOSPIRA	EQ 25MG BASE/ML	N011719 007	Mar 31, 1982
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METHOTREXATE PRESERVATIVE FREE

HOSPIRA	EQ 20MG BASE/2ML (EQ 10MG BASE/ML)	N011719 014	Apr 13, 2005
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+

	EQ 500MG BASE/20ML (EQ 25MG BASE/ML) **	N011719 013	Apr 13, 2005
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	EQ 2.5GM BASE/100ML (EQ 25MG BASE/ML)	N011719 011	Apr 13, 2005
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METHOTREXATE SODIUM

ABRAXIS PHARM

EQ 2.5MG BASE/ML	A089323 001	Jun 13, 1986
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EQ 20MG BASE/VIAL	A088935 001	Oct 11, 1985
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EQ 25MG BASE/ML	A089263 001	Jun 13, 1986
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EQ 25MG BASE/ML	A089322 001	Jun 13, 1986
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EQ 50MG BASE/VIAL	A088936 001	Oct 11, 1985
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EQ 100MG BASE/VIAL	A088937 001	Oct 11, 1985
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FRESENIUS KABI USA	EQ 250MG BASE/10ML (EQ 25MG BASE/ML)	A040263 002	Feb 26, 1999
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HOSPIRA	EQ 2.5MG BASE/ML	N011719 004	
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EQ 20MG BASE/VIAL	N011719 001	
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EQ 25MG BASE/ML	N011719 005	
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EQ 50MG BASE/VIAL	N011719 003	
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EQ 100MG BASE/VIAL	N011719 006	
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NORBROOK	EQ 25MG BASE/ML	A088648 001	May 09, 1986
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PHARMACHEMIE USA	EQ 25MG BASE/ML	A089158 001	Jul 08, 1988
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METHOTREXATE SODIUM PRESERVATIVE FREE

HOSPIRA	EQ 1GM BASE/VIAL	N011719 009	Apr 07, 1988
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MYLAN LABS LTD	EQ 50MG BASE/2ML (EQ 25MG BASE/ML)	A201529 001	Mar 29, 2012
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EQ 100MG BASE/4ML (EQ 25MG BASE/ML)	A201529 002	Mar 29, 2012
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EQ 200MG BASE/8ML (EQ 25MG BASE/ML)	A201529 003	Mar 29, 2012
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MEXATE

BRISTOL	EQ 20MG BASE/VIAL	A086358 001	
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EQ 50MG BASE/VIAL	A086358 002	
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EQ 100MG BASE/VIAL	A086358 003	
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EQ 250MG BASE/VIAL	A086358 004	
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MEXATE-AQ

BRISTOL MYERS	EQ 25MG BASE/ML	A088760 001	Feb 14, 1985
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MEXATE-AQ PRESERVED

BRISTOL MYERS SQUIBB	EQ 25MG BASE/ML	A089887 001	Apr 14, 1989
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TABLET; ORAL

METHOTREXATE SODIUM

DURAMED PHARMS BARR	EQ 2.5MG BASE	A040233 001	Jun 17, 1999
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METHOXAMINE HYDROCHLORIDE

INJECTABLE; INJECTION

VASOXYL

GLAXOSMITHKLINE	10MG/ML	N006772 002	
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20MG/ML	N006772 001	
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METHOXSALEN

CAPSULE; ORAL

8-MOP

+ VALEANT PHARM INTL	10MG	N009048 001	
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METHOXSALEN

ANI PHARMS INC	10MG	A087781 001	Jun 08, 1982
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LOTION; TOPICAL

OXSORALEN

+ VALEANT PHARM INTL	1%	N009048 002	
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METHSCOPOLAMINE BROMIDE

TABLET; ORAL

METHSCOPOLAMINE BROMIDE

BRECKENRIDGE PHARM	2.5MG	A040642 001	Dec 06, 2011
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5MG	A040642 002	Dec 06, 2011
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PVT FORM	2.5MG	A080970 001	
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VINTAGE PHARMS	2.5MG	A040624 001	Dec 28, 2006
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5MG	A040624 002	Dec 28, 2006
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PAMINE

FOUGERA PHARMS	2.5MG **	N008848 001	
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PAMINE FORTE

FOUGERA PHARMS	5MG **	N008848 002	Mar 25, 2003
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Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

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

IVAX PHARMS 2.5MG A087913 001 Jun 03, 1982

5MG A087786 001 May 18, 1982

MYLAN 2.5MG A087671 001 Aug 17, 1982

MYLAN PHARMS INC 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

YAOPHARMA CO LTD 2.5MG A089835 001 Aug 18, 1988

5MG A089837 001 Aug 18, 1988

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 A071006 001 Dec 16, 1986

500MG A071009 001 Dec 16, 1986

HALSEY 125MG A071751 001 Mar 28, 1988

250MG A071752 001 Mar 28, 1988

500MG A071753 001 Mar 28, 1988

PAR PHARM 125MG A070535 001 Jan 02, 1987

250MG A070536 001 Jan 02, 1987

500MG A070537 001 Jan 02, 1987

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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

METHYLDOPATABLET; ORAL
METHYLDOPA

	500MG	A070194	001	Apr 25, 1986
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
YAOPHARMA CO LTD	125MG	A071700	001	Mar 02, 1988
	250MG	N018934	001	Jun 29, 1984
	500MG	N018934	002	Jun 29, 1984

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

TABLET; ORAL

METHERGINE

+ EDISON THERAPS LLC	0.2MG **	N006035	003	
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METHYLPHENIDATE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

RITALIN LA

+ NOVARTIS	60MG **	N021284	005	Oct 27, 2014
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FOR SUSPENSION, EXTENDED RELEASE; ORAL

METHYLPHENIDATE HYDROCHLORIDE

ACTAVIS LABS FL INC	5MG/ML	A206049	001	May 17, 2018
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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
AUROLIFE PHARMA LLC	5MG	A209276	001	Oct 25, 2018
	10MG	A209276	002	Oct 25, 2018
	20MG	A209276	003	Oct 25, 2018
CEDIPROF INC	5MG	A208737	001	Feb 01, 2019
	10MG	A208737	002	Feb 01, 2019
	20MG	A208737	003	Feb 01, 2019
CNTY LINE PHARMS	5MG	A206840	001	Sep 15, 2016
	10MG	A206840	002	Sep 15, 2016
	20MG	A206840	003	Sep 15, 2016
LANNETT CO INC	5MG	A086429	001	
	10MG	A085799	001	
	20MG	A086428	001	

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

METHYLPHENIDATE HYDROCHLORIDE

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

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
	HERITAGE PHARMA	20MG	A075450 001	Dec 21, 2001
	PAR PHARM	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
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METHYLPREDNISOLONE

TABLET;ORAL

MEDROL

	PHARMACIA AND UPJOHN	24MG	N011153 005	
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METHYLPREDNISOLONE

	HEATHER	4MG	A085650 001	
	INVATECH	4MG	A087341 001	
	LUPIN LTD	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
	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
	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	
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INJECTABLE; INJECTION

M-PREDROL

	BEL MAR	40MG/ML	A086666 001	
		80MG/ML	A087135 001	

METHYLPREDNISOLONE ACETATE

	AKORN	40MG/ML	A086903 001	Oct 20, 1982
		80MG/ML	A086903 002	Oct 20, 1982
	AMNEAL	40MG/ML	A210043 001	May 20, 2019
		80MG/ML	A210043 002	May 20, 2019
	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	

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	

DISCONTINUED DRUG PRODUCT LIST

6-285(of 430)

** See List Footnote

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
	EQ 1GM BASE/VIAL	A089576	001	Feb 22, 1991
HOSPIRA	EQ 40MG BASE/VIAL	A040664	001	Dec 20, 2005
	EQ 40MG BASE/VIAL	A085853	001	
	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	
HOSPIRA INC	EQ 1GM BASE/VIAL	A089174	001	Aug 18, 1987
	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
	EQ 1GM BASE/VIAL	A040709	002	Feb 21, 2007
ELKINS SINN	EQ 40MG BASE/VIAL	A086906	001	
	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
INTL MEDICATION	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
TEVA PARENTERAL	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
WATSON LABS	EQ 500MG BASE/VIAL	A088523	001	Jul 24, 1984
	EQ 1GM BASE/VIAL	A088524	001	Jul 24, 1984

METHYLPREDNISOLONE; NEOMYCIN SULFATE

OINTMENT; OPHTHALMIC

NEO-MEDROL

PHARMACIA AND UPJOHN	0.1%;EQ 3.5MG BASE/GM	A060645	001	
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METHYLTESTOSTERONE

CAPSULE; ORAL

METHYLTESTOSTERONE

HEATHER	10MG	A084967	001	
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VIRILON

CHARTWELL	10MG	A087750	001	Nov 24, 1982
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TABLET; BUCCAL

ANDROID 5

VALEANT PHARM INTL	5MG	A087222	001	
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ORETON

SCHERING	10MG	A080281	001	
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Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

6-286(of 430)

** See List Footnote

METHYLTESTOSTERONE

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
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METHYLTESTOSTERONE

IMPAX LABS	25MG	A084310 001
INWOOD LABS	10MG	A080839 001
	25MG	A080973 001
KV PHARM	10MG	A084312 001
LANNETT	10MG	A087092 001
	25MG	A087111 001
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

Nov 05, 1982
Jan 27, 1983

ORETON METHYL

SCHERING	10MG	N003158 001
	25MG	N003158 002

METHYPRYLON

CAPSULE;ORAL

NOLUDAR

ROCHE	300MG	N009660 008
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ELIXIR;ORAL

NOLUDAR

ROCHE	50MG/5ML	N009660 007
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TABLET;ORAL

NOLUDAR

ROCHE	50MG	N009660 002
	200MG	N009660 004

METHYSERGIDE MALEATE

TABLET;ORAL

SANSERT

NOVARTIS	2MG	N012516 001
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METIPRANOLOL HYDROCHLORIDE

SOLUTION/DROPS;OPHTHALMIC

METIPRANOLOL

SANDOZ INC	0.3%	A075720 001	Aug 06, 2001
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OPTIPRANOLOL

+ BAUSCH AND LOMB	0.3%	N019907 001	Dec 29, 1989
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Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

METOCLOPRAMIDE HYDROCHLORIDE

CONCENTRATE; ORAL

METOCLOPRAMIDE INTENSOL

ROXANE	EQ 10MG BASE/ML	A072995 001	Jan 30, 1992
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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

WEST-WARD PHARMS INT	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
LANNETT CO INC	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
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

REGLAN

+ ROBINS AH	EQ 5MG BASE/5ML **	N018821 001	Mar 25, 1983
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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
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MAXOLON

KING PHARMS	EQ 10MG BASE	A070106 001	Mar 04, 1986
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METOCLOPRAMIDE HYDROCHLORIDE

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
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
YAOPHARMA CO LTD	EQ 5MG BASE	A074478 001	Oct 05, 1995
	EQ 10MG BASE	A072215 001	Jan 30, 1990
	EQ 10MG BASE	A074478 002	Oct 05, 1995

TABLET, ORALLY DISINTEGRATING; ORAL

METOZOLV ODT

+ SALIX PHARMS	EQ 5MG BASE	N022246 001	Sep 04, 2009
+	EQ 10MG BASE **	N022246 002	Sep 04, 2009

DISCONTINUED DRUG PRODUCT LIST

6-288(of 430)

** See List Footnote

METOCLOPRAMIDE HYDROCHLORIDETABLET, ORALLY DISINTEGRATING;ORAL
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	
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METOLAZONE

TABLET;ORAL

DIULO

GD SEARLE LLC	2.5MG	N018535 001	
	5MG	N018535 002	
	10MG	N018535 003	

METOLAZONE

ANI PHARMS INC	2.5MG	A076600 001	Jan 06, 2004
	5MG	A076833 001	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

LANNETT CO INC	0.5MG	N019532 001	Oct 30, 1987
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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

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
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
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
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METOPROLOL TARTRATE

AM REGENT	1MG/ML	A090386 001	Sep 30, 2009
LUITPOLD	1MG/ML	A091307 001	Dec 29, 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
HERITAGE PHARMA	50MG	A074141 001	Jan 31, 1995
	100MG	A074141 002	Jan 31, 1995
MYLAN	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
RENATA	50MG	A074453 001	Apr 27, 1995
	100MG	A074453 002	Apr 27, 1995
SUN PHARM INDUSTRIES	25MG	A073654 002	Jul 15, 2009
	50MG	A073654 003	Dec 21, 1993
	100MG	A073654 001	Dec 21, 1993

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

6-289(of 430)

** See List Footnote

METOPROLOL TARTRATE

TABLET; ORAL

METOPROLOL TARTRATE

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
YAOPHARMA CO LTD	50MG	A073288 001	Mar 25, 1994
	100MG	A073289 001	Mar 25, 1994

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
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INJECTABLE; INJECTION

FLAGYL I.V. RTU IN PLASTIC CONTAINER

PFIZER	500MG/100ML	N018353 002	
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METRO I.V.

B BRAUN	500MG/100ML	N018674 001	Aug 31, 1982
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METRONIDAZOLE

ABBOTT	500MG/100ML	N018889 001	Nov 18, 1983
ABRAXIS PHARM	500MG/100ML	A070071 001	Dec 03, 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
WEST-WARD PHARMS INT	500MG/100ML	N018907 001	Mar 30, 1984

TABLET; ORAL

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
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

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

METRONIDAZOLE

TABLET, EXTENDED RELEASE;ORAL

FLAGYL ER

+ GD SEARLE LLC	750MG	N020868 001	Nov 26, 1997
METRONIDAZOLE			
ABLE	750MG	A076462 001	Jun 25, 2003
ALEMBIC PHARMS LTD	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

METYRAPONE

TABLET;ORAL

METOPIRONE

HRA PHARMA	250MG	N012911 001	
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MEXILETINE HYDROCHLORIDE

CAPSULE;ORAL

MEXILETINE HYDROCHLORIDE

ANI PHARMS INC	150MG	A074450 001	May 16, 1996
	200MG	A074450 002	May 16, 1996
	250MG	A074450 003	May 16, 1996
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

MICONAZOLE

INJECTABLE;INJECTION

MONISTAT

JANSSEN PHARMA	10MG/ML	N018040 001	
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MICONAZOLE NITRATE

CREAM;TOPICAL

MONISTAT-DERM

INSIGHT PHARMS	2%	N017494 001	
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CREAM;VAGINAL

MICONAZOLE NITRATE

TEVA	2%	A074136 001	Jan 04, 1995
TEVA PHARMS	2%	A074030 001	Oct 30, 1992

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

MICONAZOLE NITRATE

CREAM, SUPPOSITORY;TOPICAL, VAGINAL

M-ZOLE 7 DUAL PACK

ACTAVIS MID ATLANTIC 2%,100MG

A074586 001 Jul 17, 1997

MICONAZOLE 7 COMBINATION PACK

ACP NIMBLE 2%,100MG

A076585 001 Mar 26, 2004

LOTION;TOPICAL

MONISTAT-DERM

INSIGHT PHARMS 2%

N017739 001

TAMPON;VAGINAL

MONISTAT 5

PERSONAL PRODS 100MG

N018592 001 Oct 27, 1989

MIDAZOLAM HYDROCHLORIDE

INJECTABLE; INJECTION

MIDAZOLAM HYDROCHLORIDE

AKORN INC EQ 5MG BASE/ML

A075481 001 Jun 30, 2000

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

HOSPIRA EQ 1MG BASE/ML

A075396 001 Jun 20, 2000

EQ 5MG BASE/ML

A075396 002 Jun 20, 2000

EQ 5MG BASE/ML

A075484 001 Jun 20, 2000

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

WOCKHARDT 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

MIDAZOLAM HYDROCHLORIDE PRESERVATIVE FREE

MYLAN ASI EQ 1MG BASE/ML

A090315 001 Nov 29, 2010

EQ 5MG BASE/ML

A090315 002 Nov 29, 2010

MIDOZALAM HYDROCHLORIDE

MYLAN ASI EQ 1MG BASE/ML

A090316 001 May 04, 2011

EQ 5MG BASE/ML

A090316 002 May 04, 2011

VERSED

+ HLR EQ 1MG BASE/ML **

N018654 002 May 26, 1987

+ EQ 5MG BASE/ML **

N018654 001 Dec 20, 1985

SYRUP;ORAL

MIDAZOLAM HYDROCHLORIDE

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

CASI PHARMS INC 2.5MG

A076514 001 Sep 11, 2003

5MG

A076514 002 Sep 11, 2003

10MG

A076514 003 Jul 02, 2004

PROAMATINE

+ SHIRE LLC 2.5MG **

N019815 001 Sep 06, 1996

+ 5MG **

N019815 002 Sep 06, 1996

+ 10MG **

N019815 003 Mar 20, 2002

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

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
USPHARMA WINDLAS	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
HOSPIRA	EQ 1MG BASE/ML	A075830 001	May 28, 2002
	EQ 1MG BASE/ML	A075884 001	May 28, 2002
MYLAN INSTITUTIONAL	EQ 1MG BASE/ML	A076428 001	Jun 16, 2003
WEST-WARD PHARMS INT	EQ 1MG BASE/ML	A075852 001	May 28, 2002
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
WEST-WARD PHARMS INT	EQ 20MG BASE/100ML (EQ 0.2MG BASE/ML)	A075510 001	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

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 67.5MG BASE	N201922 002	Jul 11, 2012
	EQ 112.5MG BASE	N201922 004	Jul 11, 2012

INJECTABLE; INJECTION

MINOCIN

LEDERLE	EQ 100MG BASE/VIAL	A062139 001	
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SUSPENSION; ORAL

MINOCIN

BAUSCH	EQ 50MG BASE/5ML	N050445 001	
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TABLET; ORAL

MINOCYCLINE HYDROCHLORIDE

+ 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 90MG BASE	A065485 002	Mar 17, 2009
	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
MYLAN	EQ 55MG BASE	A203443 001	Aug 21, 2019
MYLAN LABS LTD	EQ 65MG BASE	A201467 001	Jul 30, 2019
	EQ 115MG BASE	A201467 002	Jul 30, 2019
MYLAN PHARMS INC	EQ 45MG BASE	A090911 001	Jul 20, 2010
	EQ 90MG BASE	A090911 002	Jul 20, 2010
	EQ 135MG BASE	A090911 003	Jul 20, 2010

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

MINOCYCLINE HYDROCHLORIDETABLET, EXTENDED RELEASE;ORAL
SOLODYN

+	MEDICIS	EQ 45MG BASE **	N050808 001	May 08, 2006
+		EQ 90MG BASE **	N050808 002	May 08, 2006
+		EQ 135MG BASE **	N050808 003	May 08, 2006

MINOXIDIL

SOLUTION;TOPICAL

MINOXIDIL (FOR MEN)

	APOTEX INC	2%	A074924 001	Apr 29, 1998
	BAUSCH AND LOMB	2%	A074643 001	Apr 09, 1996
	COPLEY PHARM	2%	A074500 001	May 23, 1996
	SIGHT PHARMS	2%	A074743 002	Oct 18, 1996
	TEVA	2%	A074589 001	Apr 05, 1996

MINOXIDIL (FOR WOMEN)

	APOTEX INC	2%	A074924 002	Apr 29, 1998
	SIGHT PHARMS	2%	A074743 001	Oct 18, 1996

MINOXIDIL EXTRA STRENGTH (FOR MEN)

	APOTEX INC	5%	A075839 001	Oct 01, 2001
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TABLET;ORAL

LONITEN

+	PHARMACIA AND UPJOHN	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
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MIRABEGRON

TABLET, EXTENDED RELEASE;ORAL

MIRABEGRON

	SAWAI USA	25MG	A209446 001	Dec 27, 2019
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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
	MYLAN PHARMS INC	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 USA INC	45MG **	N020415 003	Mar 17, 1997
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Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

MIRTAZAPINETABLET, 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
	30MG	A076307 002	Dec 17, 2003
	45MG	A076307 003	Feb 28, 2006

MISOPROSTOLTABLET;ORAL
MISOPROSTOL

ANI PHARMS INC	0.1MG	A076095 001	Jul 10, 2002
	0.2MG	A076095 002	Jul 10, 2002
ZYDUS PHARMS	0.1MG	A210201 001	Jul 02, 2019
	0.2MG	A210201 002	Jul 02, 2019

MITOMYCIN

INJECTABLE; INJECTION

MITOMYCIN

HOSPIRA	20MG/VIAL	A064106 001	Nov 29, 1995
WEST-WARD PHARMS INT	5MG/VIAL	A064117 001	Apr 19, 1995

MITOZYTREX

+ SUPERGEN	5MG/VIAL **	N050763 001	Nov 14, 2002
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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
MYLAN INSTITUTIONAL	EQ 20MG BASE/10ML (EQ 2MG BASE/ML)	A078980 001	Apr 13, 2009
	EQ 30MG BASE/15ML (EQ 2MG BASE/ML)	A078980 002	Apr 13, 2009
MYLAN LABS LTD	EQ 20MG BASE/10ML (EQ 2MG BASE/ML)	A201014 001	Dec 11, 2012

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

MYLAN LABS LTD	EQ 2MG BASE/ML	A078562 001	Apr 30, 2009
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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

MODAFINIL

TABLET; ORAL

MODAFINIL

HIKMA PHARMS	100MG	A090543 001	Sep 26, 2012
	200MG	A090543 002	Sep 26, 2012

DISCONTINUED DRUG PRODUCT LIST

6-295(of 430)

** 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

+ ENDO PHARMS

5MG **

N017111 001

+

10MG **

N017111 002

+

25MG **

N017111 003

CONCENTRATE; ORAL

MOBAN

ENDO PHARMS

20MG/ML

N017938 001

TABLET; ORAL

MOBAN

+ ENDO PHARMS

5MG **

N017111 004

+

10MG **

N017111 005

+

25MG **

N017111 006

+

50MG **

N017111 007

+

100MG **

N017111 008

MOMETASONE FUROATE

CREAM; TOPICAL

ELOCON

MERCK SHARP DOHME

0.1%

N019625 001 May 06, 1987

+

0.1%

N019625 002 Apr 19, 2013

OINTMENT; TOPICAL

ELOCON

+ MERCK SHARP DOHME

0.1%

N019543 001 Apr 30, 1987

MOMETASONE FUROATE

TARO

0.1%

A076624 001 Dec 03, 2004

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

MYLAN PHARMS INC

EQ 4MG BASE/PACKET

A202776 001 Dec 18, 2012

TABLET; ORAL

MONTELUKAST SODIUM

APOTEX CORP

EQ 10MG BASE

A201294 001 Aug 03, 2012

HIKMA

EQ 10MG BASE

A090655 001 Aug 03, 2012

MYLAN PHARMS INC

EQ 10MG BASE

A079103 001 Aug 03, 2012

VINTAGE PHARMS LLC

EQ 10MG BASE

A091576 001 Aug 03, 2012

TABLET, CHEWABLE; ORAL

MONTELUKAST SODIUM

APOTEX INC

EQ 4MG BASE

A201508 001 Aug 03, 2012

EQ 5MG BASE

A201508 002 Aug 03, 2012

JUBILANT GENERICS

EQ 4MG BASE

A203795 001 Feb 27, 2015

EQ 5MG BASE

A203795 002 Feb 27, 2015

MYLAN PHARMS INC

EQ 4MG BASE

A079142 001 Aug 03, 2012

EQ 5MG BASE

A079142 002 Aug 03, 2012

VINTAGE PHARMS LLC

EQ 4MG BASE

A091588 001 Aug 03, 2012

EQ 5MG BASE

A091588 002 Aug 03, 2012

MORICIZINE HYDROCHLORIDE

TABLET; ORAL

ETHMOZINE

SHIRE

200MG

N019753 001 Jun 19, 1990

250MG

N019753 002 Jun 19, 1990

300MG

N019753 003 Jun 19, 1990

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

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

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 INC	15MG/ML	N202515 005	Nov 14, 2011
	ICU MEDICAL INC	0.5MG/ML	N019917 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;ORAL

MORPHINE SULFATE

LANNETT CO INC	10MG/5ML	A202309 001	Nov 25, 2015
	20MG/5ML	A202310 001	Oct 30, 2015
	100MG/5ML	N201517 001	Jun 23, 2011
VISTAPHARM	10MG/5ML	A201947 001	Jan 05, 2012
	20MG/5ML	A201947 002	Jan 05, 2012

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

MORPHINE SULFATE

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
MYLAN PHARMS INC	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

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

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
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TABLET;ORAL

MOXIFLOXACIN HYDROCHLORIDE

	MYLAN	EQ 400MG BASE	A204635	001	Aug 31, 2015
	SUNSHINE LAKE	EQ 400MG BASE	A206295	001	Sep 28, 2018

MUPIROCIN

OINTMENT;TOPICAL

BACTROBAN

+	GLAXOSMITHKLINE	2% **	N050591	001	Dec 31, 1987
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MUPIROCIN CALCIUM

CREAM;TOPICAL

BACTROBAN

+	GLAXOSMITHKLINE	EQ 2% BASE **	N050746	001	Dec 11, 1997
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OINTMENT;NASAL

BACTROBAN

+	GLAXOSMITHKLINE	EQ 2% BASE	N050703	001	Sep 18, 1995
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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
	ZYDUS PHARMS USA INC	250MG	A065433	001	May 04, 2009

TABLET;ORAL

MYCOPHENOLATE MOFETIL

	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
	OXFORD PHARMS	500MG	A090606	001	Jul 16, 2010
	TEVA PHARMS	500MG	A065457	001	May 04, 2009
	ZYDUS PHARMS USA INC	500MG	A065477	001	May 04, 2009

NABUMETONE

TABLET;ORAL

NABUMETONE

	COPLEY PHARM	750MG	A075179	001	Jun 06, 2000
	OXFORD PHARMS	500MG	A079093	001	Feb 27, 2009
		750MG	A079093	002	Feb 27, 2009
	SANDOZ	500MG	A075590	001	Feb 25, 2002
		750MG	A075590	002	Feb 25, 2002

RELAFEN

+	SMITHKLINE BEECHAM	500MG **	N019583	001	Dec 24, 1991
+		750MG **	N019583	002	Dec 24, 1991

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

NADOLOL

TABLET; ORAL

CORCARD

US WORLDMEDS LLC	120MG	N018063 003	
	160MG	N018063 004	

NADOLOL

HERITAGE PHARMA	120MG	A074255 002	Jan 24, 1996
	160MG	A074255 003	Jan 24, 1996
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	
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FOR SOLUTION; ORAL

UNIPEN

WYETH AYERST	EQ 250MG BASE/5ML	N050199 001	
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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	
MYLAN LABS LTD	EQ 1GM BASE/VIAL	A200002 001	Apr 07, 2014
	EQ 2GM BASE/VIAL	A200002 002	Apr 07, 2014
SANDOZ	EQ 500MG BASE/VIAL	A062527 001	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	
+	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
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TABLET; ORAL

UNIPEN

WYETH AYERST	EQ 500MG BASE	N050462 001	
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NAFTIFINE HYDROCHLORIDE

CREAM; TOPICAL

NAFTIN

+	SEBELA IRELAND LTD	1%	N019599 001	Feb 29, 1988
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GEL; TOPICAL

NAFTIFINE HYDROCHLORIDE

TARO	2%	A208201 001	Apr 10, 2019
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NALBUPHINE HYDROCHLORIDE

INJECTABLE; INJECTION

NALBUPHINE

ABRAXIS PHARM	10MG/ML	A070751 001	Jul 02, 1986
	20MG/ML	A070752 001	Sep 24, 1986

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

NALBUPHINE HYDROCHLORIDE

INJECTABLE; INJECTION

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
MYLAN LABS LTD	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			
+ PAR PHARM INC	10MG/ML **	N018024 001	
+	20MG/ML **	N018024 002	May 27, 1982

NALIDIXIC ACID

SUSPENSION; ORAL

NEGRAM

SANOFI AVENTIS US	250MG/5ML	N017430 001	
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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

NEGRAM

SANOFI AVENTIS US	250MG	N014214 002	
	500MG	N014214 004	
	1GM	N014214 005	

NALMEFENE HYDROCHLORIDE

INJECTABLE; INJECTION

REVEX

+ EUROHLTH INTL SARL	EQ 0.1MG BASE/ML **	N020459 001	Apr 17, 1995
+	EQ 1MG BASE/ML **	N020459 002	Apr 17, 1995

NALOXONE HYDROCHLORIDE

INJECTABLE; INJECTION

NALOXONE

WEST-WARD PHARMS INT	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
	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
EUROHLTH INTL SARL	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	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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

NALOXONE HYDROCHLORIDE

INJECTABLE; INJECTION

NALOXONE HYDROCHLORIDE

	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
	1MG/ML	A072115 001	Apr 27, 1988
MARSAM PHARMS LLC	0.4MG/ML	A071811 001	Jul 19, 1988
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
VIRTUS PHARMS	0.4MG/ML	A207846 001	Dec 17, 2018
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
SPRAY, METERED; NASAL			
NALOXONE HYDROCHLORIDE			
TEVA PHARMS USA	4MG/SPRAY	A209522 001	Apr 19, 2019
NARCAN			
+ ADAPT	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

NALOXONE HYDROCHLORIDE; PENTAZOCINE HYDROCHLORIDE

TABLET; ORAL

TALWIN NX

SANOFI AVENTIS US	EQ 0.5MG BASE; EQ 50MG BASE **	N018733 001	Dec 16, 1982
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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 INC	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

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

NANDROLONE DECANOATE

INJECTABLE; INJECTION

DECA-DURABOLIN

ASPEN GLOBAL INC	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
AKORN	100MG/ML	A087519 001	Sep 28, 1983
AM REGENT	200MG/ML	A091252 001	Aug 30, 2010
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			
AKORN INC	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

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
FOSUN PHARMA	250MG	A074140 001	Dec 21, 1993
	375MG	A074140 002	Dec 21, 1993
	500MG	A074140 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
IVAX SUB TEVA PHARMS	250MG	A074111 001	Feb 28, 1995
	375MG	A074111 002	Feb 28, 1995
	500MG	A074111 003	Feb 28, 1995
MYLAN	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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

NAPROXEN

TABLET;ORAL
NAPROXEN

	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
FOSUN PHARMA	375MG	A075061 001	Feb 18, 1998
	500MG	A075061 002	Feb 18, 1998
MYLAN PHARMS INC	375MG	A075390 001	Apr 19, 2001
	500MG	A075390 002	Apr 19, 2001

NAPROXEN SODIUM

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
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
MYLAN	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
TEVA	EQ 250MG BASE	A074142 001	Dec 21, 1993
	EQ 500MG BASE	A074142 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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

NAPROXEN SODIUM; SUMATRIPTAN SUCCINATE

TABLET; ORAL

TREXIMET

+ CURRAX

60MG;EQ 10MG BASE

N021926 002 May 14, 2015

NARATRIPTAN HYDROCHLORIDE

TABLET; ORAL

NARATRIPTAN

ANI PHARMS INC

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

CASI PHARMS INC

EQ 1MG BASE

A090288 001 Jul 07, 2010

EQ 2.5MG BASE

A090288 002 Jul 07, 2010

MYLAN PHARMS INC

EQ 1MG BASE

A202431 001 May 31, 2012

EQ 2.5MG BASE

A202431 002 May 31, 2012

NATEGLINIDE

TABLET; ORAL

NATEGLINIDE

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

AMERIGEN PHARMS LTD

EQ 2.5MG BASE

A203659 001 Apr 16, 2015

EQ 5MG BASE

A203659 002 Apr 16, 2015

EQ 10MG BASE

A203659 003 Apr 16, 2015

EQ 20MG BASE

A203659 004 Apr 16, 2015

GLENMARK PHARMS LTD

EQ 2.5MG BASE

A203821 001 May 25, 2017

EQ 5MG BASE

A203821 002 May 25, 2017

EQ 10MG BASE

A203821 003 May 25, 2017

EQ 20MG BASE

A203821 004 May 25, 2017

INDCHEMIE HEALTH

EQ 2.5MG BASE

A203828 001 Jul 29, 2015

EQ 5MG BASE

A203828 002 Jul 29, 2015

EQ 10MG BASE

A203828 003 Jul 29, 2015

EQ 20MG BASE

A203828 004 Jul 29, 2015

TORRENT

EQ 2.5MG BASE

A203966 001 Mar 02, 2018

EQ 5MG BASE

A203966 002 Mar 02, 2018

EQ 10MG BASE

A203966 003 Mar 02, 2018

EQ 20MG BASE

A203966 004 Mar 02, 2018

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

+ ALLERGAN

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

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

NEFAZODONE HYDROCHLORIDE

TABLET; ORAL

NEFAZODONE HYDROCHLORIDE

ANI PHARMS INC	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
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
FOSUN PHARMA	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
IVAX SUB TEVA PHARMS	50MG	A075763 001	Sep 16, 2003
	100MG	A075763 002	Sep 16, 2003
	150MG	A075763 003	Sep 16, 2003
	200MG	A075763 004	Sep 16, 2003
	250MG	A075763 005	Sep 16, 2003
MYLAN	100MG	A076129 002	Sep 16, 2003
	150MG	A076129 003	Sep 16, 2003
	200MG	A076129 004	Sep 16, 2003
	250MG	A076129 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
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NEOMYCIN SULFATE

SOLUTION; ORAL

MYCIFRADIN

PHARMACIA AND UPJOHN	EQ 87.5MG BASE/5ML	N050285 001	
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NEO-FRADIN

X GEN PHARMS	EQ 87.5MG BASE/5ML	A065010 001	May 23, 2002
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TABLET; ORAL

MYCIFRADIN

PHARMACIA AND UPJOHN	EQ 350MG BASE	A060520 001	
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NEOBIOTIC

PFIZER	EQ 350MG BASE	A060475 001	
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NEOMYCIN SULFATE

BRISTOL MYERS SQUIBB	500MG	A060365 001	
LANNETT	500MG	A060607 001	
LANNETT CO INC	500MG	A204435 001	Jun 10, 2016
LILLY	500MG	A060385 001	

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

6-305(of 430)

** See List Footnote

NEOMYCIN SULFATE

TABLET; ORAL

NEOMYCIN SULFATE

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
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OINTMENT; OPHTHALMIC

STATROL

ALCON	EQ 3.5MG BASE/GM;10,000 UNITS/GM	N050344	002
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SOLUTION; IRRIGATION

NEOMYCIN AND POLYMYXIN B SULFATE

WATSON LABS	EQ 40MG BASE/ML;200,000 UNITS/ML	A062664	001	Apr 08, 1986
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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	

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
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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
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NEOMYCIN SULFATE; PREDNISOLONE SODIUM PHOSPHATE

OINTMENT; OPHTHALMIC

NEO-HYDELTRASOL

MERCK	EQ 3.5MG BASE/GM;EQ 0.25% PHOSPHATE	N050378	001
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NEOMYCIN SULFATE; TRIAMCINOLONE ACETONIDE

CREAM; TOPICAL

MYTRES A

SAVAGE LABS	EQ 3.5MG BASE/GM;0.1%	A062598	001	Jul 21, 1986
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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
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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

NESIRITIDE RECOMBINANT

FOR SOLUTION; INTRAVENOUS

NATRECOR

+ SCIOS LLC	1.5MG/VIAL	N020920	001	Aug 10, 2001
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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

TABLET; ORAL

NEVIRAPINE

APOTEX INC	200MG	A203021	001	May 22, 2012
TECH ORGANIZED	200MG	A203176	001	May 22, 2012

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

NEVIRAPINE

TABLET, EXTENDED RELEASE;ORAL

NEVIRAPINE

APOTEX	400MG	A205258 001	Apr 03, 2014
CIPLA	400MG	A206448 001	Oct 15, 2015
MYLAN	100MG	A206271 001	Nov 09, 2015
TECH ORGANIZED	100MG	A207467 001	Jul 31, 2017
	400MG	A207467 002	Jul 31, 2017

NIACIN

CAPSULE;ORAL

WAMPOCAP

MEDPOINTE PHARM HLC	500MG	N011073 003	
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TABLET;ORAL

NIACIN

EVERYLIFE	500MG	A083203 001	
HALSEY	500MG	A083453 001	
HIKMA PHARMS	500MG	A083718 001	
IMPAX LABS	500MG	A083115 001	
IVAX SUB TEVA PHARMS	500MG	A083180 001	
MK LABS	500MG	A083525 001	
PUREPAC PHARM	500MG	A083271 001	
SANDOZ	500MG	A083306 001	
TABLICAPS	500MG	A084237 001	
WATSON LABS	500MG	A083136 001	
	500MG	A083305 001	
	500MG	A085172 001	

NICOLAR

SANOFI AVENTIS US	500MG	A083823 001	
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TABLET, EXTENDED RELEASE;ORAL

NIACIN

MYLAN	500MG	A203742 001	Feb 22, 2019
	750MG	A203742 002	Feb 22, 2019
	1GM	A203742 003	Feb 22, 2019
YICHANG HUMANWELL	500MG	A212017 001	Jun 10, 2019
	750MG	A212017 002	Jun 10, 2019
	1GM	A212017 003	Jun 10, 2019

NIASPAN

ABBVIE	375MG	N020381 001	Jul 28, 1997
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NIASPAN TITRATION STARTER PACK

ABBVIE	375MG;500MG;750MG	N020381 005	Jul 28, 1997
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NIACINAMIDE; PYRIDOXINE HYDROCHLORIDE; TYROSINE

SUSPENSION;ORAL

TPN

INTL MINERALS	15MG/5ML;3.75MG/5ML;600MG/5ML	N008378 003	
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NICARDIPINE HYDROCHLORIDE

CAPSULE;ORAL

CARDENE

CHIESI USA INC	20MG **	N019488 001	Dec 21, 1988
	30MG **	N019488 002	Dec 21, 1988

NICARDIPINE HYDROCHLORIDE

ANI PHARMS INC	20MG	A074439 001	Dec 10, 1996
	20MG	A074540 001	Oct 28, 1996
	20MG	A074670 001	Oct 28, 1996
	30MG	A074439 002	Dec 10, 1996
	30MG	A074540 002	Oct 28, 1996
	30MG	A074670 002	Oct 28, 1996

CAPSULE, EXTENDED RELEASE;ORAL

CARDENE SR

+ CHIESI USA INC	30MG **	N020005 001	Feb 21, 1992
+ CHIESI USA INC	45MG **	N020005 002	Feb 21, 1992
+ CHIESI USA INC	60MG **	N020005 003	Feb 21, 1992

INJECTABLE; INJECTION

CARDENE

+ CHIESI USA INC	25MG/10ML (2.5MG/ML) **	N019734 001	Jan 30, 1992
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NICARDIPINE HYDROCHLORIDE

AM REGENT	25MG/10ML (2.5MG/ML)	A090534 001	Nov 17, 2009
MYLAN INSTITUTIONAL	25MG/10ML (2.5MG/ML)	A090664 001	Nov 17, 2009
NAVINTA LLC	25MG/10ML (2.5MG/ML)	A090125 001	Nov 17, 2009

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

6-307(of 430)

** See List Footnote

NICARDIPINE HYDROCHLORIDE

INJECTABLE; INJECTION

NICARDIPINE HYDROCHLORIDE

SUN PHARMA GLOBAL	25MG/10ML (2.5MG/ML)	N078405	001	Nov 17, 2009
WEST-WARD PHARMS INT	25MG/10ML (2.5MG/ML)	A078714	001	Dec 28, 2009
WOCKHARDT	25MG/10ML (2.5MG/ML)	A090671	001	Nov 17, 2009

INJECTABLE; INTRAVENOUS

CARDENE IN 5.0% DEXTROSE IN PLASTIC CONTAINER

+ CHIESI USA INC	40MG/200ML (0.2MG/ML)	N019734	005	Nov 07, 2008
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NICARDIPINE HYDROCHLORIDE IN 0.9% SODIUM CHLORIDE

EXELA PHARMA SCIENCE	20MG/200ML (0.1MG/ML)	N022276	002	Apr 07, 2016
	40MG/200ML (0.2MG/ML)	N022276	003	Apr 07, 2016

NICLOSAMIDE

TABLET, CHEWABLE; ORAL

NICLOCIDE

BAYER PHARMS	500MG	N018669	001	May 14, 1982
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NICOTINE

FILM, EXTENDED RELEASE; TRANSDERMAL

NICOTROL

MCNEIL CONS	15MG/16HR	N020536	001	Jul 03, 1996
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PROSTEP

AVEVA	11MG/24HR	N019983	003	Dec 23, 1998
	22MG/24HR	N019983	004	Dec 23, 1998

NICOTINE POLACRILEX

GUM, CHEWING; BUCCAL

NICOTINE POLACRILEX

IVAX SUB TEVA PHARMS	EQ 2MG BASE	A076880	001	Feb 18, 2009
	EQ 4MG BASE	A077850	001	Feb 18, 2009

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

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
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TABLET, EXTENDED RELEASE; ORAL

AFEDITAB CR

WATSON LABS	60MG	A075659	001	Oct 26, 2001
WATSON LABS TEVA	30MG	A075128	001	Mar 10, 2000

NIFEDIPINE

MARTEC USA LLC	90MG	A075414	003	Mar 23, 2004
MYLAN	30MG	A075108	001	Dec 17, 1999
	30MG	A090649	001	Jun 21, 2010
	60MG	A090649	002	Jun 21, 2010
	90MG	A090649	003	Jun 21, 2010
MYLAN LABS LTD	30MG	A090602	001	Sep 13, 2010
	60MG	A090602	002	Sep 13, 2010
	90MG	A090602	003	Sep 13, 2010

NILUTAMIDE

TABLET; ORAL

NILANDRON

CONCORDIA	50MG	N020169	001	Sep 19, 1996
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NIMODIPINE

CAPSULE; ORAL

NIMODIPINE

SUN PHARM INDS INC	30MG	A077067	001	Apr 17, 2007
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NIMOTOP

+ BAYER PHARMS	30MG **	N018869	001	Dec 28, 1988
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Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

NISOLDIPINETABLET, EXTENDED RELEASE;ORAL
SULAR

+	COVIS PHARMA BV	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

NITRIC OXIDE

GAS;INHALATION

INOMAX

+	MALLINCKRODT HOSP	100PPM **	N020845 002	Dec 23, 1999
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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

	LANNETT	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	
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NITROFURANTOIN, MACROCRYSTALLINE

CAPSULE;ORAL

NITROFURANTOIN

	INVATECH	25MG	A074336 001	Jan 25, 1995
		50MG	A074336 002	Jan 25, 1995
		100MG	A074336 003	Jan 25, 1995
	MYLAN	50MG	A074967 001	Jul 09, 1997
		100MG	A074967 002	Jul 09, 1997
		100MG	A077025 001	Aug 18, 2004
	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)

	RANBAXY LABS LTD	75MG;25MG	A076951 001	Mar 30, 2005
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NITROFUZAZONE

CREAM;TOPICAL

FURACIN

	SHIRE	0.2%	A083789 001	
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DRESSING;TOPICAL

ACTIN-N

	SHERWOOD MEDCL	0.2%	N017343 001	
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Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

NITROFURAZONE

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

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

MEDICIS 0.4MG/HR A089773 001 Aug 30, 1996

VALEANT PHARMS 0.1MG/HR A089771 001 Aug 30, 1996

VALEANT PHARMS 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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

NITROGLYCERIN

INJECTABLE; INJECTION

TRIDIL

HOSPIRA	0.5MG/ML	N018537 002	Jun 16, 1983
	5MG/ML	N018537 001	

NIZATIDINE

CAPSULE; ORAL

AXID

SMITHKLINE BEECHAM	150MG	N019508 001	Apr 12, 1988
	300MG	N019508 002	Apr 12, 1988

NIZATIDINE

ANI PHARMS INC	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

SOLUTION; ORAL

AXID

+ BRAINTREE	15MG/ML **	N021494 001	May 25, 2004
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NONOXYNOL-9

AEROSOL; VAGINAL

DELFIN

PERSONAL PRODS	12.5%	N014349 002	
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NOREPINEPHRINE BITARTRATE

INJECTABLE; INJECTION

NOREPINEPHRINE BITARTRATE

METRICS PHARM	EQ 1MG BASE/ML	A040522 001	Sep 30, 2004
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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	
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NORETHINDRONE

TABLET; ORAL

NORLUTIN

PARKE DAVIS	5MG	N010895 002	
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TABLET; ORAL-28

MICRONOR

+ JANSSEN PHARMS	0.35MG	N016954 001	
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NORETHINDRONE ACETATE

TABLET; ORAL

AYGESTIN

+ DURAMED RES	5MG **	N018405 001	Apr 21, 1982
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NORETHINDRONE ACETATE

AUROBINDO PHARMA LTD	5MG	A204236 001	Jan 08, 2016
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NORLUTATE

PARKE DAVIS	5MG	N012184 002	
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NORFLOXACIN

SOLUTION/DROPS; OPHTHALMIC

CHIBROXIN

MERCK	0.3%	N019757 001	Jun 17, 1991
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TABLET; ORAL

NOROXIN

+ MERCK	400MG **	N019384 002	Oct 31, 1986
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NORGESTREL

TABLET; ORAL

OPILL

+ LABORATOIRE HRA	0.075MG	N017031 001	
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DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

NORTRIPTYLINE HYDROCHLORIDE

CAPSULE; ORAL

AVENTYL HYDROCHLORIDE

LILLY

EQ 10MG BASE

N014684 001

EQ 25MG BASE

N014684 002

NORTRIPTYLINE HYDROCHLORIDE

ANI PHARMS INC

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

AUROLIFE PHARMA LLC

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

MYLAN

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

SOLUTION; ORAL

AVENTYL

+ RANBAXY

EQ 10MG BASE/5ML **

N014685 001

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

TARO

100,000 UNITS/GM

A062457 001 Jul 28, 1983

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

+ DAVA PHARMS INC

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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

NYSTATIN

SUSPENSION; ORAL

MYCOSTATIN

DELCOR ASSET CORP 100,000 UNITS/ML A061533 001

NILSTAT

+ GLENMARK GENERICS 100,000 UNITS/ML ** N050299 001

NYSTATIN

ACP NIMBLE 100,000 UNITS/ML A062776 001 Dec 17, 1987

ALPHARMA US PHARMS 100,000 UNITS/ML A062571 001 Oct 29, 1985

G AND W LABS INC 100,000 UNITS/ML A062349 001 Jul 14, 1982

MORTON GROVE 100,000 UNITS/ML A062835 001 Nov 19, 1987

PHARMADERM 100,000 UNITS/ML A062518 001 Jul 06, 1984

PHARMAFAIR 100,000 UNITS/ML A062541 001 Jan 16, 1985

SOCORRO 100,000 UNITS/ML A062832 001 Dec 27, 1991

TEVA 100,000 UNITS/ML A062670 001 Jun 18, 1987

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

QUANTUM PHARMICS 500,000 UNITS A062525 001 Oct 29, 1984

SANDOZ 500,000 UNITS A062065 001

UPSHER SMITH LABS 500,000 UNITS A062524 001 Nov 26, 1985

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

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

DELCOR 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

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

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

MYTRES F

SAVAGE LABS 100,000 UNITS/GM; 0.1% A062601 001 Oct 09, 1985

NYSTATIN AND TRIAMCINOLONE ACETONIDE

CROWN LABS INC 100,000 UNITS/GM; 0.1% A207731 001 Dec 26, 2017

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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

NYSTATIN; TRIAMCINOLONE ACETONIDE

OINTMENT; TOPICAL

NYSTATIN AND TRIAMCINOLONE ACETONIDE

ZYDUS PHARMS	100,000 UNITS/GM;0.1%	A204764	001	Nov 08, 2018
NYSTATIN-TRIAMCINOLONE ACETONIDE				
PHARMADERM	100,000 UNITS/GM;0.1%	A062603	001	Oct 09, 1985

OCTREOTIDE ACETATE

INJECTABLE; INJECTION

OCTREOTIDE ACETATE

HERITAGE PHARMS INC	EQ 0.05MG BASE/ML	A204669	001	Dec 27, 2018
	EQ 0.1MG BASE/ML	A204669	002	Dec 27, 2018
	EQ 0.2MG BASE/ML	A203765	001	Sep 07, 2018
	EQ 0.5MG BASE/ML	A204669	003	Dec 27, 2018
	EQ 1MG BASE/ML	A203765	002	Sep 07, 2018
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
	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

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
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SOLUTION/DROPS; OPHTHALMIC

OFLOXACIN

SANDOZ	0.3%	A076848	001	Nov 25, 2008
SANDOZ INC	0.3%	A076231	001	May 14, 2004

SOLUTION/DROPS; OTIC

FLOXIN OTIC

+ DAIICHI	0.3% **	N020799	001	Dec 16, 1997
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OFLOXACIN

ALVOGEN	0.3%	A090395	001	Aug 11, 2009
SANDOZ INC	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

LARKEN LABS	200MG	A076093	001	Sep 02, 2003
	300MG	A076093	002	Sep 02, 2003
	400MG	A076093	003	Sep 02, 2003
RANBAXY LABS LTD	200MG	A076220	001	Sep 02, 2003
	300MG	A076220	002	Sep 02, 2003
	400MG	A076220	003	Sep 02, 2003

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

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
MYLAN	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
SUNSHINE LAKE	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

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
HEC PHARM	5MG	A208146 001	Jul 02, 2018
	10MG	A208146 002	Jul 02, 2018
	15MG	A208146 003	Jul 02, 2018
	20MG	A208146 004	Jul 02, 2018

OLMESARTAN MEDOXOMIL

TABLET; ORAL

OLMESARTAN MEDOXOMIL

JUBILANT GENERICS	5MG	A205482 001	Apr 24, 2017
	20MG	A205482 002	Apr 24, 2017
	40MG	A205482 003	Apr 24, 2017
TEVA PHARMS USA	5MG	A091079 001	Apr 24, 2017
	20MG	A091079 002	Apr 24, 2017
	40MG	A091079 003	Apr 24, 2017

OLOPATADINE HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

OLOPATADINE HYDROCHLORIDE

ZAMBON SPA	EQ 0.1% BASE	A204706 001	Dec 07, 2015
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OMBITASVIR; PARITAPREVIR; RITONAVIR

TABLET; ORAL

TECHNIVIE

+ ABBVIE INC	12.5MG; 75MG; 50MG	N207931 001	Jul 24, 2015
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OMEGA-3-ACID ETHYL ESTERS

CAPSULE; ORAL

OMEGA-3-ACID ETHYL ESTERS

PAR PHARM INC	1GM CONTAINS AT LEAST 900MG OF THE ETHYL ESTERS OF OMEGA-3 FATTY ACIDS	A091018 001	Jun 24, 2014
ZYDUS	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	1.2GM CONTAINS AT LEAST 900MG OF THE ETHYL ESTERS OF OMEGA-3 FATTY ACIDS	N204977 001	Apr 23, 2014
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Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

OMEGA-3-CARBOXYLIC ACIDS

CAPSULE;ORAL

EPANOVA

+	ASTRAZENECA PHARMS	1GM CONTAINS AT LEAST 850MG OF POLYUNSATURATED FATTY ACIDS	N205060 001	May 05, 2014
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OMEPRAZOLE

CAPSULE, DELAYED REL PELLETS;ORAL

OMEPRAZOLE

LUPIN LTD	40MG	A202384 001	Aug 25, 2015
MYLAN	10MG	A075876 001	May 29, 2003
	10MG	A205070 001	Jun 29, 2018
	20MG	A075876 002	May 29, 2003
	20MG	A205070 002	Jun 29, 2018
	40MG	A075876 003	Jan 21, 2009
	40MG	A205070 003	Jun 29, 2018

PRILOSEC

+	ASTRAZENECA PHARMS	10MG **	N019810 003	Oct 05, 1995
+		20MG **	N019810 001	Sep 14, 1989
+		40MG **	N019810 002	Jan 15, 1998

TABLET, DELAYED RELEASE;ORAL

OMEPRAZOLE

APOTEX	20MG	A210070 001	Feb 11, 2019
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OMEPRAZOLE; SODIUM BICARBONATE

CAPSULE;ORAL

OMEPRAZOLE AND SODIUM BICARBONATE

PAR PHARM	20MG;1.1GM	A078966 001	May 25, 2010
	40MG;1.1GM	A078966 002	May 25, 2010

ONDANSETRON

TABLET, ORALLY DISINTEGRATING;ORAL

ONDANSETRON

CHARTWELL MOLECULES	4MG	A077406 003	Dec 26, 2006
	8MG	A077406 004	Dec 26, 2006
	16MG	A077406 001	Dec 26, 2006
	24MG	A077406 002	Dec 26, 2006
NESHER PHARMS	4MG	A077717 001	Jun 25, 2007
	8MG	A077717 002	Jun 25, 2007

ONDANSETRON HYDROCHLORIDE

INJECTABLE;INJECTION

ONDANSETRON HYDROCHLORIDE

APOTEX INC	EQ 2MG BASE/ML	A077368 001	Dec 26, 2006
HOSPIRA	EQ 2MG BASE/ML	A076695 001	Dec 26, 2006
LANNETT CO INC	EQ 2MG BASE/ML	A090116 001	Apr 14, 2010
	EQ 2MG BASE/ML	A090883 001	Aug 05, 2010
LUITPOLD	EQ 2MG BASE/ML	A077582 001	Dec 26, 2006
	EQ 2MG BASE/ML	A079039 001	Nov 18, 2008
MYLAN LABS LTD	EQ 2MG BASE/ML	A078257 001	Apr 23, 2008
PLIVA HRVATSKA DOO	EQ 2MG BASE/ML	A077544 001	Dec 26, 2006
SAGENT PHARMS	EQ 2MG BASE/ML	A078180 001	Mar 26, 2007
SUN PHARM INDS (IN)	EQ 2MG BASE/ML	A077172 001	Dec 26, 2006

ONDANSETRON HYDROCHLORIDE AND DEXTROSE IN PLASTIC CONTAINER

HOSPIRA	EQ 0.64MG BASE/ML	A076978 001	Feb 26, 2007
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ONDANSETRON HYDROCHLORIDE PRESERVATIVE FREE

AM REGENT	EQ 2MG BASE/ML	A079032 001	Nov 18, 2008
APOTEX INC	EQ 2MG BASE/ML	A077343 001	Dec 26, 2006
HIKMA FARMACEUTICA	EQ 2MG BASE/ML	A076780 001	Dec 26, 2006
HOSPIRA	EQ 2MG BASE/ML	A076696 001	Dec 26, 2006
LUITPOLD	EQ 2MG BASE/ML	A077387 001	Dec 26, 2006
MYLAN LABS LTD	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

+	NOVARTIS	EQ 2MG BASE/ML **	N020007 001	Jan 04, 1991
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ZOFRAN AND DEXTROSE IN PLASTIC CONTAINER

+	GLAXOSMITHKLINE	EQ 0.64MG BASE/ML **	N020403 001	Jan 31, 1995
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ZOFRAN PRESERVATIVE FREE

+	NOVARTIS	EQ 2MG BASE/ML **	N020007 003	Dec 10, 1993
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Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

ONDANSETRON HYDROCHLORIDE

TABLET; ORAL

ONDANSETRON HYDROCHLORIDE

CHARTWELL MOLECULES	EQ 4MG BASE	A077303 001	Jun 25, 2007
	EQ 8MG BASE	A077303 002	Jun 25, 2007
	EQ 24MG BASE	A077303 004	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
MYLAN	EQ 4MG BASE	A076930 001	Jun 25, 2007
	EQ 8MG BASE	A076930 002	Jun 25, 2007
	EQ 24MG BASE	A076930 004	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

ORPHENADRINE CITRATE

INJECTABLE; INJECTION

NORFLEX

+ TELIGENT	30MG/ML	N013055 001	
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ORPHENADRINE CITRATE

WATSON LABS	30MG/ML	A087062 001	
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TABLET, EXTENDED RELEASE; ORAL

NORFLEX

+ MEDICIS	100MG	N012157 001	
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ORPHENADRINE CITRATE

ASCOT	100MG	A088067 001	Apr 06, 1983
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SANDOZ	100MG	A085046 001	
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WATSON LABS	100MG	A084303 001	
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ORPHENADRINE HYDROCHLORIDE

TABLET; ORAL

DISIPAL

3M	50MG	N010653 001	
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OSELTAMIVIR PHOSPHATE

FOR SUSPENSION; ORAL

TAMIFLU

ROCHE	EQ 12MG BASE/ML	N021246 001	Dec 14, 2000
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OXACILLIN SODIUM

CAPSULE; ORAL

BACTOCILL

GLAXOSMITHKLINE	EQ 250MG BASE	A061336 001	
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	EQ 250MG BASE	A062241 001	
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	EQ 500MG BASE	A061336 002	
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	EQ 500MG BASE	A062241 002	
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OXACILLIN SODIUM

ANI PHARMS INC	EQ 250MG BASE	A062222 001	
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	EQ 500MG BASE	A062222 002	
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APOTHECON	EQ 250MG BASE	A061450 002	
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	EQ 500MG BASE	A061450 001	
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PROSTAPHLIN

APOTHECON	EQ 500MG BASE	N050118 002	
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FOR SOLUTION; ORAL

BACTOCILL

GLAXOSMITHKLINE	EQ 250MG BASE/5ML	A062321 001	
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OXACILLIN SODIUM

APOTHECON	EQ 250MG BASE/5ML	A061457 001	
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TEVA	EQ 250MG BASE/5ML	A062252 001	
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PROSTAPHLIN

APOTHECON	EQ 250MG BASE/5ML	N050194 001	
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INJECTABLE; INJECTION

BACTOCILL

GLAXOSMITHKLINE	EQ 500MG BASE/VIAL **	A061334 009	Mar 26, 1982
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	EQ 1GM BASE/VIAL **	A061334 006	Mar 26, 1982
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	EQ 1GM BASE/VIAL **	A062736 001	Dec 19, 1986
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	EQ 2GM BASE/VIAL **	A061334 007	Mar 26, 1982
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	EQ 2GM BASE/VIAL **	A062736 002	Dec 19, 1986
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	EQ 4GM BASE/VIAL **	A061334 008	Mar 26, 1982
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	EQ 10GM BASE/VIAL **	A061334 010	
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Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

OXACILLIN SODIUM

INJECTABLE; INJECTION

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 INC	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
	MYLAN LABS LTD	EQ 1GM BASE/VIAL	A091486 001	Aug 25, 2014
		EQ 2GM BASE/VIAL	A091486 002	Aug 25, 2014
	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
	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
+		100MG/VIAL **	N021492 002	Aug 09, 2002
+		200MG/40ML (5MG/ML) **	N021759 003	Nov 17, 2006

OXALIPLATIN

AM REGENT

	50MG/10ML (5MG/ML)	A204378 001	May 12, 2017
	100MG/20ML (5MG/ML)	A204378 002	May 12, 2017

FRESENIUS KABI ONCOL

	50MG/VIAL	A078810 001	Aug 07, 2009
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HOSPIRA INC

	100MG/VIAL	A078810 002	Aug 07, 2009
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HOSPIRA INC

	50MG/VIAL	A078815 001	Sep 30, 2009
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HOSPIRA INC

	100MG/VIAL	A078815 002	Sep 30, 2009
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MYLAN LABS LTD

	50MG/VIAL	A200979 001	Aug 08, 2012
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MYLAN LABS LTD

	100MG/VIAL	A200979 002	Aug 08, 2012
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MYLAN LABS LTD

	200MG/40ML (5MG/ML)	A091358 003	Nov 14, 2017
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SANDOZ

	50MG/VIAL	A090849 001	Apr 28, 2011
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SANDOZ

	100MG/VIAL	A090849 002	Apr 28, 2011
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SANDOZ INC

	50MG/10ML (5MG/ML)	A078812 001	Aug 07, 2009
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SANDOZ INC

	100MG/20ML (5MG/ML)	A078812 002	Aug 07, 2009
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SUN PHARM

	50MG/VIAL	A078818 001	Aug 07, 2009
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SUN PHARM

	50MG/10ML (5MG/ML)	A202922 001	Apr 08, 2014
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SUN PHARM

	100MG/VIAL	A078818 002	Aug 07, 2009
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SUN PHARM

	100MG/20ML (5MG/ML)	A202922 002	Apr 08, 2014
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SUN PHARM

	200MG/40ML (5MG/ML)	A202922 003	Feb 15, 2019
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Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

OXAMNIQUINE

CAPSULE; ORAL

VANSIL

PFIZER

250MG

N018069 001

OXANDROLONE

TABLET; ORAL

OXANDRIN

+ GEMINI LABS LLC

2.5MG

N013718 001

+

10MG

N013718 002 Nov 05, 2001

OXANDROLONE

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

OXAPROZIN

TABLET; ORAL

OXAPROZIN

ACTAVIS ELIZABETH

600MG

A075843 001 Oct 03, 2001

BEXIMCO PHARMS USA

600MG

A075842 001 Apr 12, 2001

MYLAN

600MG

A075851 001 Aug 17, 2001

MYLAN PHARMS INC

600MG

A075847 001 Feb 28, 2001

SANDOZ

600MG

A075850 001 Apr 27, 2001

SUN PHARM INDS INC

600MG

A075844 001 Jan 03, 2002

WATSON LABS

600MG

A075848 001 Feb 09, 2001

OXAPROZIN POTASSIUM

TABLET; ORAL

DAYPRO ALTA

GD SEARLE

600MG

N020776 001 Oct 17, 2002

OXAZEPAM

CAPSULE; ORAL

OXAZEPAM

AM THERAP

10MG

A071955 001 Mar 03, 1988

15MG

A071956 001 Mar 03, 1988

30MG

A071957 001 Mar 03, 1988

FRONTIDA BIOPHARM

10MG

A071026 002 Aug 10, 1987

15MG

A071026 003 Aug 10, 1987

30MG

A071026 001 Aug 10, 1987

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

DISCONTINUED DRUG PRODUCT LIST

6-319(of 430)

** See List Footnote

OXCARBAZEPINE

TABLET; ORAL

OXCARBAZEPINE

ANI PHARMS INC	150MG	A078005 001	Dec 11, 2007
	300MG	A078005 002	Dec 11, 2007
	600MG	A078005 003	Dec 11, 2007
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

OXPRENOLOL 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
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OXTRIPHYLLINE

MORTON GROVE	100MG/5ML	A088243 001	Dec 05, 1983
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SYRUP; ORAL

CHOLEDYL

PARKE DAVIS	50MG/5ML	N009268 011	
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OXTRIPHYLLINE PEDIATRIC

MORTON GROVE	50MG/5ML	A088242 001	Dec 05, 1983
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TABLET, DELAYED RELEASE; ORAL

CHOLEDYL

PARKE DAVIS	100MG	N009268 003	
	200MG	N009268 007	

OXTRIPHYLLINE

WATSON LABS	100MG	A087866 001	Aug 25, 1983
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	200MG	A087835 001	Aug 25, 1983
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TABLET, EXTENDED RELEASE; ORAL

CHOLEDYL SA

WARNER CHILCOTT LLC	400MG	A087863 001	May 24, 1983
	600MG	A086742 001	

OXYBUTYNIN

FILM, EXTENDED RELEASE; TRANSDERMAL

OXYBUTYNIN

BARR LABS DIV TEVA	3.9MG/24HR	A090526 001	Mar 04, 2014
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GEL, METERED; TRANSDERMAL

GELNIQUE 3%

+ ALLERGAN	3%	N202513 001	Dec 07, 2011
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OXYBUTYNIN CHLORIDE

GEL; TRANSDERMAL

OXYBUTYNIN CHLORIDE

PAR PHARM INC	10% (100MG/PACKET)	A207329 001	May 31, 2018
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SYRUP; ORAL

DITROPAN

+ ORTHO MCNEIL JANSSEN	5MG/5ML **	N018211 001	
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OXYBUTYNIN CHLORIDE

ANDA REPOSITORY	5MG/5ML	A075039 001	Jan 29, 1999
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LANNETT CO INC	5MG/5ML	A076682 001	Dec 28, 2004
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PHARM ASSOC	5MG/5ML	A074997 001	Oct 15, 1997
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TABLET; ORAL

DITROPAN

+ JANSSEN PHARMS	5MG **	N017577 001	
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OXYBUTYNIN CHLORIDE

QUANTUM PHARMICS	5MG	A072296 001	Dec 08, 1988
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USL PHARMA	5MG	A070746 001	Mar 10, 1988
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WATSON LABS	5MG	A072485 001	Apr 19, 1989
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Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

6-320(of 430)

** See List Footnote

OXYBUTYNIN CHLORIDE

TABLET, EXTENDED RELEASE;ORAL

DITROPAN XL

+	JANSSEN PHARMS	15MG **	N020897	003	Jun 22, 1999
	OXYBUTYNIN CHLORIDE				
	MYLAN	5MG	A076702	001	Nov 09, 2006
	MYLAN PHARMS INC	10MG	A076644	001	Nov 09, 2006
		15MG	A076644	002	May 10, 2007

OXYCODONE HYDROCHLORIDE

SOLUTION;ORAL

OXYCODONE HYDROCHLORIDE

	ANI PHARMS INC	100MG/5ML	A203447	001	Aug 30, 2017
	HIKMA	100MG/5ML	A203208	001	Jul 12, 2013
	LANNETT CO INC	100MG/5ML	A204085	001	Sep 09, 2014

TABLET;ORAL

OXYCODONE HYDROCHLORIDE

	VINTAGE PHARMS	5MG	A077712	003	Mar 02, 2009
		10MG	A077712	004	Apr 13, 2015
		15MG	A077712	001	Jan 31, 2007
		20MG	A077712	005	Apr 13, 2015
		30MG	A077712	002	Jan 31, 2007

ROXYBOND

	INSPIRION DELIVERY	5MG	N209777	001	Apr 20, 2017
		15MG	N209777	002	Apr 20, 2017
		30MG	N209777	003	Apr 20, 2017

TABLET, EXTENDED RELEASE;ORAL

ROXICODONE

	ROXANE	10MG	N020932	001	Oct 26, 1998
		30MG	N020932	002	Oct 26, 1998

OXYMETAZOLINE HYDROCHLORIDE

SOLUTION/DROPS;OPHTHALMIC

OCUCLEAR

	BAYER HEALTHCARE LLC	0.025%	N018471	001	May 30, 1986
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OXYMORPHONE HYDROCHLORIDE

INJECTABLE;INJECTION

OPANA

+	ENDO PHARMS	1MG/ML	N011707	002	
		1.5MG/ML	N011707	001	

SUPPOSITORY;RECTAL

NUMORPHAN

	ENDO PHARMS	5MG	N011738	004	
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TABLET;ORAL

OPANA

+	ENDO PHARMS	5MG	N021611	001	Jun 22, 2006
+		10MG	N021611	002	Jun 22, 2006

TABLET, EXTENDED RELEASE;ORAL

OPANA ER

	ENDO PHARMS	5MG **	N021610	001	Jun 22, 2006
+		5MG	N201655	001	Dec 09, 2011
		7.5MG **	N021610	005	Feb 29, 2008
+		7.5MG	N201655	002	Dec 09, 2011
		10MG **	N021610	002	Jun 22, 2006
+		10MG	N201655	003	Dec 09, 2011
		15MG **	N021610	006	Feb 29, 2008
+		15MG	N201655	004	Dec 09, 2011
		20MG **	N021610	003	Jun 22, 2006
+		20MG	N201655	005	Dec 09, 2011
		30MG **	N021610	007	Feb 29, 2008
+		30MG	N201655	006	Dec 09, 2011
		40MG **	N021610	004	Jun 22, 2006
+		40MG	N201655	007	Dec 09, 2011

OXYMORPHONE HYDROCHLORIDE

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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

OXYMORPHONE HYDROCHLORIDETABLET, EXTENDED RELEASE;ORAL
OXYMORPHONE HYDROCHLORIDE

	40MG	A200792 007	Oct 24, 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

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 PHARMS 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

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

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		

PACLITAXEL

INJECTABLE; INJECTION

PACLITAXEL					
ACCORD HLTHCARE	6MG/ML	A075436	001	Nov 12, 2004	
	6MG/ML	A205720	001	Aug 17, 2018	
HOSPIRA	6MG/ML	A076233	001	Aug 01, 2002	
MYLAN	6MG/ML	A075278	001	Jan 25, 2002	
PLIVA LACHEMA	6MG/ML	A077413	001	Mar 12, 2008	
SANDOZ INC	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	

PALIPERIDONE

TABLET, EXTENDED RELEASE; ORAL

INVEGA					
+ JANSSEN PHARMS	12MG **	N021999	004	Dec 19, 2006	

PALONOSETRON HYDROCHLORIDE

CAPSULE; ORAL

ALOXI					
+ HELSINN HLTHCARE	EQ 0.5MG BASE **	N022233	001	Aug 22, 2008	

INJECTABLE; INTRAVENOUS

PALONOSETRON HYDROCHLORIDE					
DR REDDYS LABS LTD	EQ 0.075MG BASE/1.5ML (EQ 0.05MG BASE/ML)	A201533	001	Apr 21, 2016	

SOLUTION; INTRAVENOUS

PALONOSETRON HYDROCHLORIDE					
DR REDDYS LABS LTD	EQ 0.075MG BASE/1.5ML (EQ 0.05MG BASE/ML)	N203050	001	Mar 01, 2016	
	EQ 0.25MG BASE/5ML (EQ 0.05MG BASE/ML)	N203050	002	Mar 01, 2016	

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	
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	

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

PANCRELIPASE (AMYLASE;LIPASE;PROTEASE)

CAPSULE;ORAL

COTAZYM

ORGANON USA INC	30,000USP UNITS;8,000USP UNITS;30,000USP UNITS	N020580 001	Dec 09, 1996
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CAPSULE, DELAYED RELEASE;ORAL

ULTRESA

+ FOREST LABS INC	27,600USP UNITS;13,800USP UNITS;27,600USP UNITS	N022222 001	Mar 01, 2012
+	41,400USP UNITS;20,700USP UNITS;41,400USP UNITS	N022222 002	Mar 01, 2012
+	46,000USP UNITS;23,000USP UNITS;46,000USP UNITS	N022222 003	Mar 01, 2012

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	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	

PANTOPRAZOLE SODIUM

INJECTABLE; IV (INFUSION)

PANTOPRAZOLE SODIUM

MYLAN LABS LTD	EQ 40MG BASE/VIAL	A208580 001	May 04, 2018
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TABLET, DELAYED RELEASE;ORAL

PANTOPRAZOLE SODIUM

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

PARAMETHADIONE

CAPSULE;ORAL

PARADIONE

ABBVIE	150MG	N006800 003	
	300MG	N006800 001	

SOLUTION;ORAL

PARADIONE

ABBVIE	300MG/ML	N006800 002	
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PARAMETHASONE ACETATE

TABLET;ORAL

HALDRONE

LILLY	1MG	N012772 005	
	2MG	N012772 006	

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
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Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

PAROMOMYCIN SULFATE

CAPSULE; ORAL

HUMATIN

KING PFIZER	EQ 250MG BASE	A062310	001
PARKE DALE	EQ 250MG BASE	A060521	001

SYRUP; ORAL

HUMATIN

PARKE DAVIS	EQ 125MG BASE/5ML	A060522	001
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PAROXETINE HYDROCHLORIDE

CAPSULE; ORAL

PAXIL

+	APOTEX TECHNOLOGIES	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
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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
MYLAN PHARMS INC	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
	EQ 40MG BASE	A075566	004	Mar 08, 2004
PAXIL				
APOTEX TECHNOLOGIES	EQ 50MG BASE	N020031	004	Dec 29, 1992

PAZOPANIB HYDROCHLORIDE

TABLET; ORAL

VOTRIENT

+	NOVARTIS	EQ 400MG BASE	N022465	002	Oct 19, 2009
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PEGADEMASE BOVINE

INJECTABLE; INJECTION

ADAGEN

+	LEADIANT BIOSCI INC	250 UNITS/ML	N019818	001	Mar 21, 1990
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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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

6-325(of 430)

** See List Footnote

PEMIROLAST POTASSIUM

SOLUTION/DROPS;OPHTHALMIC

ALAMAST

SANTEN 0.1% N021079 001 Sep 24, 1999

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

+ AUXILIUM PHARMS LLC 10MG ** N018976 001 Dec 30, 1987
 + 20MG ** N018976 004 Jan 05, 1989

PENICILLAMINE

CAPSULE;ORAL

CUPRIMINE

VALEANT PHARMS INTL 125MG N019853 002

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 efficacy 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
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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

AM ANTIBIOTICS	EQ 125MG BASE/5ML		A061529 001
	EQ 250MG BASE/5ML		A061529 002
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

PFIZERPEN VK

PFIZER	EQ 125MG BASE/5ML		A061815 001
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DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PENICILLIN V POTASSIUM

FOR SOLUTION;ORAL

PFIZERPEN VK

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

AM ANTIBIOTICS

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

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

6-329(of 430)

** 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 YB-169

INJECTABLE; INJECTION

YTTERBIUM YB 169 DTPA

3M 2mCi/ML N017518 001

PENTOBARBITAL

ELIXIR; ORAL

NEMBUTAL

OAK PHARMS 18.2MG/5ML A083244 001

PENTOBARBITAL SODIUM

CAPSULE; ORAL

NEMBUTAL SODIUM

OAK PHARMS 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

INJECTABLE; INJECTION

PENTOBARBITAL SODIUM

ELKINS SINN 50MG/ML A083270 001

SODIUM PENTOBARBITAL

WYETH AYERST 50MG/ML A083261 001

SUPPOSITORY; RECTAL

NEMBUTAL

OAK PHARMS 30MG A083247 001 Jan 25, 1982

60MG A083247 002 Jan 25, 1982

120MG A083247 003 Jan 25, 1982

200MG A083247 004 Jan 25, 1982

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

PENTOBARBITAL SODIUM

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				
MYLAN INSTITUTIONAL	10MG/VIAL		A203554	001 Sep 19, 2014

PENTOXIFYLLINE

TABLET, EXTENDED RELEASE; ORAL

PENTOXIFYLLINE				
ANI PHARMS INC	400MG		A074878	001 Jul 09, 1997
	400MG		A075107	001 Sep 04, 1998
	400MG		A075199	001 Sep 03, 1999
HERITAGE PHARMS INC	400MG		A074877	001 Jul 08, 1997
IMPAX LABS	400MG		A075093	001 Aug 10, 1999
MYLAN	400MG		A074425	001 Jul 08, 1997
PLIVA	400MG		A074874	001 May 25, 1999
TRENTAL				
+ US PHARM HOLDINGS	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	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
PAR PHARM	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
PERINDOPRIL ERBUMINE				
ANI PHARMS INC	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
LUPIN LTD	2MG		A078263	001 Jan 27, 2010
	4MG		A078263	002 Jan 27, 2010
	8MG		A078263	003 Jan 27, 2010

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

PERMETHRIN

CREAM; TOPICAL

ELIMITE

+ MYLAN

5%

N019855 001 Aug 25, 1989

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 INC

2MG

A089707 001 Sep 10, 1987

4MG

A089708 001 Sep 10, 1987

8MG

A089456 001 Sep 10, 1987

16MG

A089457 001 Sep 10, 1987

TRILAFON

+ SCHERING

2MG **

N010775 001

+

4MG **

N010775 002

+

8MG **

N010775 003

+

16MG **

N010775 004

TABLET, EXTENDED RELEASE; ORAL

TRILAFON

SCHERING

8MG

N011361 002

PHENACEMIDE

TABLET; ORAL

PHENURONE

+ ABBVIE

500MG **

N007707 001

PHENAZOPYRIDINE HYDROCHLORIDE; SULFAMETHOXAZOLE

TABLET; ORAL

AZO GANTANOL

+ ROCHE

100MG; 500MG **

N013294 001 Sep 10, 1987

PHENAZOPYRIDINE HYDROCHLORIDE; SULFAMETHOXAZOLE; TRIMETHOPRIM

TABLET; ORAL

SULFAMETHOXAZOLE AND TRIMETHOPRIM AND PHENAZOPYRIDINE HYDROCHLORIDE

ABLE

200MG, N/A, N/A; N/A, 800MG, 160MG

N021105 001 Jun 26, 2001

PHENAZOPYRIDINE HYDROCHLORIDE; SULFISOXAZOLE

TABLET; ORAL

AZO GANTRISIN

+ ROCHE

50MG; 500MG **

N019358 001 Aug 31, 1990

PHENDIMETRAZINE TARTRATE

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 PHARMS

35MG

A085695 001

VITARINE

35MG

A085634 001

35MG

A085645 001

35MG

A085670 001

35MG

A086403 001

35MG

A086408 001

35MG

A086410 001

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PHENDIMETRAZINE TARTRATE

CAPSULE; ORAL

PHENDIMETRAZINE TARTRATE

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 PHARMS

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

FERNDALE 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

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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

6-333(of 430)

** See List Footnote

PHENDIMETRAZINE TARTRATE

TABLET;ORAL

PHENDIMETRAZINE TARTRATE

	35MG	A085525 001
MFG CHEMISTS	35MG	A085914 001
NEXGEN PHARMA INC	35MG	A086020 001
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 PHARMS	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
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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

DIBENZYLINE

+ CONCORDIA	10MG	N008708 001
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PHENPROCOUMON

TABLET;ORAL

LIQUAMAR

ORGANON USA INC	3MG	N011228 001
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PHENSUXIMIDE

CAPSULE;ORAL

MILONTIN

PARKE DAVIS	500MG	N008855 004
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DISCONTINUED DRUG PRODUCT LIST

6-334(of 430)

** See List Footnote

PHENTERMINE HYDROCHLORIDE

CAPSULE; ORAL

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
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
IVAX PHARMS	30MG	A086329 001	
LANNETT CO INC	30MG	A091359 001	Jul 16, 2010
SANDOZ	30MG	A087208 001	
	30MG	A087223 001	
	37.5MG	A088414 001	Oct 19, 1983
SUN PHARM INDUSTRIES	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	
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PHENTERMINE HYDROCHLORIDE

ABLE	37.5MG	A040402 001	Aug 30, 2001
ACTAVIS ELIZABETH	37.5MG	A040276 001	Nov 25, 1998
CHARTWELL RX	8MG	A083923 001	
	8MG	A085319 001	
	37.5MG	A087805 001	Dec 06, 1982
	37.5MG	A088596 001	Apr 04, 1984
IVAX PHARMS	8MG	A085553 001	
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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

6-335(of 430)

** See List Footnote

PHENTERMINE HYDROCHLORIDE

TABLET;ORAL

PHENTERMINE HYDROCHLORIDE	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

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			
FOSUN PHARMA	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; PROMETHAZINE HYDROCHLORIDE

SYRUP;ORAL

PHENERGAN VC			
+ ANI PHARMS	5MG/5ML;6.25MG/5ML **	N008604 003	Apr 02, 1984
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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

PHENYLEPHRINE HYDROCHLORIDE; PROMETHAZINE HYDROCHLORIDE

SYRUP;ORAL

PROMETHAZINE VC PLAIN

CENCI	5MG/5ML;6.25MG/5ML	A088815 001	Nov 22, 1985
WOCKHARDT	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	
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PHENYTOIN

SUSPENSION;ORAL

DILANTIN-30

PARKE DAVIS	30MG/5ML	N008762 002	
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PHENYTOIN

ACTAVIS MID ATLANTIC	125MG/5ML	A089892 001	Sep 25, 1992
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PHENYTOIN SODIUM

CAPSULE;ORAL

DIPHENYLAN SODIUM

LANNETT	30MG PROMPT	A080857 001	
	100MG PROMPT	A080857 002	

EXTENDED PHENYTOIN SODIUM

ANI PHARMS INC	100MG EXTENDED	A040435 001	Jun 20, 2003
	100MG EXTENDED	A089441 001	Dec 18, 1986
SUN PHARM INDS (IN)	100MG EXTENDED	A040621 001	Dec 11, 2006
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
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PHENYTOIN SODIUM

PHARMERAL	100MG PROMPT	A085435 001	
WATSON LABS	100MG PROMPT	A085894 001	

PROMPT PHENYTOIN SODIUM

ANI PHARMS INC	100MG PROMPT	A080259 001	
WATSON LABS	100MG PROMPT	A080905 001	

INJECTABLE;INJECTION

DILANTIN

PARKE DAVIS	50MG/ML	N010151 001	
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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

+ TELIGENT	1MG/0.5ML **	N012223 002	
+	10MG/ML **	N012223 001	

KONAKION

ROCHE	1MG/0.5ML	N011745 001	
	10MG/ML	N011745 003	

PHYTONADIONE

GLAXOSMITHKLINE	1MG/0.5ML	A084060 001	
	10MG/ML	A084060 002	

VITAMIN K1

HOSPIRA	10MG/ML	A087956 001	Jul 25, 1983
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DISCONTINUED DRUG PRODUCT LIST

6-337(of 430)

** See List Footnote

PILOCARPINE

INSERT, EXTENDED RELEASE;OPHTHALMIC

OCUSERT PILO-20

AKORN

5MG

N017431 001

OCUSERT PILO-40

AKORN

11MG

N017548 001

PILOCARPINE HYDROCHLORIDE

GEL;OPHTHALMIC

PILOPINE HS

ALCON

4%

N018796 001 Oct 01, 1984

TABLET;ORAL

PILOCARPINE HYDROCHLORIDE

AUROBINDO PHARMA LTD

5MG

A212377 001 Aug 13, 2019

7.5MG

A212377 002 Aug 13, 2019

PIMAVANSERIN TARTRATE

TABLET;ORAL

NUPLAZID

+

ACADIA PHARMS INC

EQ 17MG BASE

N207318 001 Apr 29, 2016

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

ACP NIMBLE

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

MYLAN PHARMS INC

EQ 15MG BASE

A076801 001 Aug 17, 2012

EQ 30MG BASE

A076801 002 Aug 17, 2012

EQ 45MG BASE

A076801 003 Aug 17, 2012

PIPECURONIUM BROMIDE

INJECTABLE; INJECTION

ARDUAN

ORGANON USA INC

10MG/VIAL

N019638 001 Jun 26, 1990

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

PIPERACETAZINETABLET; 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
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

PIPERAZINE CITRATE

SYRUP; ORAL

ANTEPAR

GLAXOSMITHKLINE	EQ 500MG BASE/5ML	N009102 001
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BRYREL

SANOFI AVENTIS US	EQ 500MG BASE/5ML	N017796 001
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MULTIFUGE

BLULINE	EQ 500MG BASE/5ML	N009452 001
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PIPERAZINE CITRATE

ALPHARMA US PHARMS	EQ 500MG BASE/5ML	A080774 001
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LANNETT	EQ 500MG BASE/5ML	A080963 001
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LUITPOLD	EQ 500MG BASE/5ML	A080671 001
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VERMIDOL

SOLVAY	EQ 500MG BASE/5ML	A080992 001
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TABLET; ORAL

ANTEPAR

GLAXOSMITHKLINE	EQ 500MG BASE	N009102 003
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PIPERAZINE CITRATE

IMPAX LABS	EQ 250MG BASE	A080874 001
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PIPERONYL BUTOXIDE; PYRETHRINS

AEROSOL; TOPICAL

RID MOUSSE

BAYER HEALTHCARE LLC	4%;EQ 0.33% BASE	N021043 001	Mar 07, 2000
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PIPOBROMAN

TABLET; ORAL

VERCYTE

ABBOTT	10MG	N016245 001
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	25MG	N016245 002
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PIRBUTEROL ACETATE

AEROSOL, METERED; INHALATION

MAXAIR

BAUSCH	EQ 0.2MG BASE/INH	N019009 001	Dec 30, 1986
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MEDICIS	EQ 0.2MG BASE/INH	N020014 001	Nov 30, 1992
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PIRFENIDONE

TABLET; ORAL

ESBRIET

+	GENENTECH INC	534MG **	N208780 002	Jan 11, 2017
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DISCONTINUED DRUG PRODUCT LIST

6-339(of 430)

** See List Footnote

PIROXICAM

CAPSULE; ORAL

PIROXICAM

BRECKENRIDGE	10MG	A208991 001	Feb 21, 2018
	20MG	A208991 002	Feb 21, 2018
CYCLE PHARMS LTD	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
IVAX SUB TEVA PHARMS	10MG	A074148 001	Jun 03, 1996
	20MG	A074148 002	Jun 03, 1996
MYLAN	10MG	A074043 001	Sep 22, 1992
	10MG	A074102 001	Jul 31, 1992
	20MG	A074043 002	Sep 22, 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 CALCIUM

TABLET; ORAL

PITAVASTATIN CALCIUM

MYLAN	EQ 1MG BASE	A206070 001	Apr 04, 2019
	EQ 2MG BASE	A206070 002	Apr 04, 2019
	EQ 4MG BASE	A206070 003	Apr 04, 2019

PITAVASTATIN MAGNESIUM

TABLET; ORAL

ZYPITAMAG

+ MEDICURE

EQ 1MG BASE N208379 001 Jul 14, 2017

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

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

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

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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

POLYETHYLENE GLYCOL 3350; POTASSIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE

FOR SOLUTION;ORAL

PEG-3350, POTASSIUM CHLORIDE, SODIUM BICARBONATE, SODIUM CHLORIDE

MYLAN	420GM/BOT;1.48GM/BOT;5.72GM/BOT;11.2GM/BOT	A090409	001	Apr 02, 2010
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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
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SOLUTION;ORAL

OCL

HOSPIRA	6GM/100ML;75MG/100ML;168MG/100ML;146MG/100ML;1.29GM/100ML	N019284	001	Apr 30, 1986
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POLYETHYLENE GLYCOL 3350; POTASSIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE; SODIUM SULFATE ANHYDROUS

FOR SOLUTION;ORAL

COLYTE

MYLAN SPECIALITY LP	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-FLAVORED

MYLAN SPECIALITY LP	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
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PEG 3350 AND ELECTROLYTES

MYLAN

	236GM/BOT;2.97GM/BOT;6.74GM/BOT;5.86GM/BOT;22.74GM/BOT	A090928	001	Jan 28, 2010
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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
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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
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COLOVAGE

DYNAPHARM

	227.1GM/PACKET;2.82GM/PACKET;6.36GM/PACKET;5.53GM/PACKET;21.5GM/PACKET	A071320	001	Apr 20, 1988
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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
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GLYCOPREP

GOLDLINE

	236GM/BOT;2.97GM/BOT;6.74GM/BOT;5.86GM/BOT;22.74GM/BOT	A072319	001	Dec 23, 1988
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GO-EVAC

VINTAGE PHARMS

	236GM/BOT;2.97GM/BOT;6.74GM/BOT;5.86GM/BOT;22.74GM/BOT	A073433	001	Apr 28, 1992
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PEG-LYTE

SANDOZ

	236GM/BOT;2.97GM/BOT;6.74GM/BOT;5.86GM/BOT;22.74GM/BOT	A073098	001	Aug 31, 1993
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POLYMYXIN B SULFATE

INJECTABLE; INJECTION

AEROSPORIN

GLAXOSMITHKLINE

	EQ 500,000 UNITS BASE/VIAL	A062036	001	
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POLYMYXIN B SULFATE

WEST-WARD PHARMS INT

	EQ 500,000 UNITS BASE/VIAL	A060716	001	
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POWDER; FOR RX COMPOUNDING

POLY-RX

X GEN PHARMS

	100,000,000 UNITS/BOT	A061578	001	
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DISCONTINUED DRUG PRODUCT LIST

6-341(of 430)

** See List Footnote

POLYMYXIN B SULFATE

POWDER; FOR RX COMPOUNDING

POLYMYXIN B SULFATE

PADDOCK LLC

100,000,000 UNITS/BOT

A062455 001 Jul 27, 1983

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

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

POTASSIUM CHLORIDE

NESHER PHARMS

10MEQ

A070980 001 Feb 17, 1987

TEVA

8MEQ

A073531 001 Apr 26, 1996

10MEQ

A073532 001 Apr 26, 1996

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

AKORN

2MEQ/ML

A088286 001 Sep 05, 1985

BAXTER HLTHCARE

2MEQ/ML

A080203 001

2MEQ/ML

A085499 001

FRESENIUS KABI USA

2MEQ/ML

A087817 001 Oct 20, 1982

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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

POTASSIUM CHLORIDE

INJECTABLE; INJECTION

POTASSIUM CHLORIDE

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
TABLET, EXTENDED RELEASE; ORAL				
K+10				
FUTURE PAK	10MEQ	A070999	001	Oct 22, 1987
K+8				
FUTURE PAK	8MEQ	A070998	001	Jan 25, 1993
KAON CL				
SAVAGE LABS	6.7MEQ	N017046	001	
KAON CL-10				
SAVAGE LABS	10MEQ	N017046	002	
KLOTRIX				
APOTHECON	10MEQ	N017850	001	
POTASSIUM CHLORIDE				
COPLEY PHARM	8MEQ	A070618	001	Sep 09, 1987
NESHER PHARMS	20MEQ	A076044	001	Apr 05, 2002
PII	10MEQ	A206630	001	Mar 29, 2019
	15MEQ	A206630	002	Mar 29, 2019
	20MEQ	A206630	003	Mar 29, 2019
+ SCHERING	10MEQ **	N019439	002	Jun 13, 1986
+	20MEQ **	N019439	001	Jun 13, 1986
STRIDES PHARMA	8MEQ	A206881	001	Jan 22, 2019
	10MEQ	A206881	002	Jan 22, 2019
	10MEQ	A210097	001	Jun 17, 2019
	20MEQ	A210098	001	Apr 26, 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% IN PLASTIC CONTAINER

B BRAUN 75MG/100ML; 900MG/100ML N018722 001 Nov 09, 1982

BAXTER HLTHCARE 75MG/100ML; 900MG/100ML N017648 004

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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

POTASSIUM CHLORIDE; SODIUM CHLORIDE

INJECTABLE; INJECTION

SODIUM CHLORIDE 0.9% AND POTASSIUM CHLORIDE 0.3% IN PLASTIC CONTAINER

B BRAUN

300MG/100ML; 900MG/100ML

N018722 004 Nov 09, 1982

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

POTASSIUM IODIDE

SOLUTION; ORAL

POTASSIUM IODIDE

ROXANE

1GM/ML **

N018551 001 Feb 19, 1982

TABLET; ORAL

THYRO-BLOCK

MEDA PHARMS

130MG

N018307 001

POTASSIUM PERCHLORATE

CAPSULE; ORAL

PERCHLORACAP

MALLINCKRODT

200MG

N017551 001

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

PRALIDOXIME CHLORIDE

INJECTABLE; INJECTION

PRALIDOXIME CHLORIDE

BAXTER HLTHCARE CORP

300MG/ML

N018799 001 Dec 13, 1982

TABLET; ORAL

PROTOPAM CHLORIDE

WYETH AYERST

500MG

N014122 002

PRAMIPEXOLE DIHYDROCHLORIDE

TABLET; ORAL

MIRAPEX

BOEHRINGER INGELHEIM

1.25MG

N020667 004 Jul 01, 1997

PRAMIPEXOLE DIHYDROCHLORIDE

HERITAGE PHARMA

0.125MG

A078551 001 Oct 08, 2010

0.125MG

A090241 001 Oct 08, 2010

0.125MG

A091254 001 Nov 30, 2010

0.25MG

A078551 002 Oct 08, 2010

0.25MG

A090241 002 Oct 08, 2010

0.25MG

A091254 002 Nov 30, 2010

0.5MG

A078551 003 Oct 08, 2010

0.5MG

A090241 003 Oct 08, 2010

0.5MG

A091254 003 Nov 30, 2010

0.75MG

A090241 004 Oct 08, 2010

0.75MG

A091254 004 Nov 30, 2010

1MG

A078551 004 Oct 08, 2010

1MG

A090241 005 Oct 08, 2010

1MG

A091254 005 Nov 30, 2010

1.5MG

A078551 005 Oct 08, 2010

1.5MG

A090241 006 Oct 08, 2010

1.5MG

A091254 006 Nov 30, 2010

MYLAN

0.125MG

A077854 001 Oct 08, 2010

0.25MG

A077854 002 Oct 08, 2010

0.5MG

A077854 003 Oct 08, 2010

0.75MG

A090764 001 Apr 09, 2010

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

6-344(of 430)

** See List Footnote

PRAMIPEXOLE DIHYDROCHLORIDE

TABLET; ORAL

PRAMIPEXOLE DIHYDROCHLORIDE

	1MG	A077854 004	Oct 08, 2010
	1.5MG	A077854 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

PRAMLINTIDE ACETATE

INJECTABLE; SUBCUTANEOUS

SYMLIN

ASTRAZENECA AB	EQ 3MG BASE/5ML (EQ 600MCG BASE/ML)	N021332 001	Mar 16, 2005
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PRAVASTATIN SODIUM

TABLET; ORAL

PRAVACHOL

+ BRISTOL MYERS SQUIBB 10MG **

N019898 002 Oct 31, 1991

PRAVASTATIN SODIUM

MYLAN

	10MG	A077013 001	Oct 23, 2006
	20MG	A077013 002	Oct 23, 2006
	40MG	A077013 003	Oct 23, 2006
	80MG	A077013 004	Dec 28, 2007
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

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

TABLET; ORAL

CENTRAX

PARKE DAVIS

10MG

N017415 001

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 INC

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

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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

6-345(of 430)

** See List Footnote

PRAZOSIN HYDROCHLORIDE

CAPSULE;ORAL

PRAZOSIN HYDROCHLORIDE

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

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

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

A040571 001 Aug 25, 2005

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

FERNISOLONE-P

FERNDALE LABS 5MG

A083941 001

PREDNISOLONE

AUROLIFE PHARMA LLC 5MG

A084773 001

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

LANNETT 5MG

A080531 002

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

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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

PREDNISOLONE

TABLET; ORAL

PREDNISOLONE

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
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PREDNISOLONE ACETATE

INJECTABLE; INJECTION

METICORTELONE

SCHERING	25MG/ML	N010255	002
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PREDNISOLONE ACETATE

AKORN	25MG/ML	A083032	001
	50MG/ML	A084492	001
BEL MAR	25MG/ML	A083738	001
	50MG/ML	A083738	002
CENT PHARMS	25MG/ML	A084717	001
	50MG/ML	A084717	002
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
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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

EYEVANCE	0.125%	N017468	001
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PREDNISOLONE ACETATE; SULFACETAMIDE SODIUM

OINTMENT; OPHTHALMIC

CETAPRED

ALCON	0.25%;10%	A087771	001	Aug 06, 1993
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METIMYD

SCHERING	0.5%;10%	N010210	002	Sep 09, 1984
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PREDSULFAIR

PHARMAFAIR	0.5%;10%	A088032	001	Apr 15, 1983
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VASOCIDIN

NOVARTIS	0.5%;10%	A088791	001	Oct 05, 1984
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SUSPENSION; OPHTHALMIC

ISOPTO CETAPRED

ALCON	0.25%;10%	A087547	001
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SUSPENSION/DROPS; OPHTHALMIC

METIMYD

SCHERING	0.5%;10%	N010210	001
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PREDAMIDE

AKORN	0.5%;10%	A088059	001	Jul 29, 1983
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PREDSULFAIR

PHARMAFAIR	0.5%;10%	A088007	001	Apr 19, 1983
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PREDSULFAIR II

PHARMAFAIR	0.2%;10%	A088837	001	Dec 24, 1985
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SULPHRIN

BAUSCH AND LOMB	0.5%;10%	A088089	001	Dec 28, 1982
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PREDNISOLONE SODIUM PHOSPHATE

INJECTABLE; INJECTION

HYDELTRASOL

MERCK	EQ 20MG PHOSPHATE/ML	N011583	002
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PREDNISOLONE SODIUM PHOSPHATE

WATSON LABS	EQ 20MG PHOSPHATE/ML	A080517	001
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Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

6-347(of 430)

** See List Footnote

PREDNISOLONE SODIUM PHOSPHATE

OINTMENT;OPHTHALMIC, OTIC

HYDELTRASOL

MERCK

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

MEDICIS PHARMS

EQ 15MG BASE/5ML

A075250 001 Jul 12, 2002

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

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

AKORN

EQ 0.11% PHOSPHATE

A083358 001

EQ 0.9% PHOSPHATE

A083358 002

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

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

PREDNISOLONE SODIUM PHOSPHATE; SULFACETAMIDE SODIUM

SOLUTION/DROPS;OPHTHALMIC

SULFACETAMIDE SODIUM AND PREDNISOLONE SODIUM PHOSPHATE

SANDOZ INC

EQ 0.23% PHOSPHATE;10%

A073630 001 May 27, 1993

SULSTER

AKORN

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

PREDNISONE

SOLUTION;ORAL

PREDNISONE

WOCKHARDT

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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

6-348(of 430)

** See List Footnote

PREDNISON

TABLET;ORAL

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

AUROLIFE PHARMA LLC 5MG A084774 001

10MG A089983 001 Jan 12, 1989

20MG A085813 001

50MG A089984 001 Jan 12, 1989

BUNDY 5MG A083676 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

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

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

LANNETT 5MG A080514 001

20MG A084275 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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

6-349(of 430)

** See List Footnote

PREDNISON

TABLET; ORAL

PREDNISON

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	
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
	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	5MG	A080223	001	

PREGABALIN

CAPSULE; ORAL

PREGABALIN

MYLAN	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
	225MG	A091228	007	Sep 20, 2019
	300MG	A091228	008	Sep 20, 2019

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 efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

6-350(of 430)

** See List Footnote

PRIMIDONE

SUSPENSION; ORAL

MYSOLINE

NURO PHARMA 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
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

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 INC	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

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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

6-351(of 430)

** See List Footnote

PROCAINAMIDE HYDROCHLORIDE

INJECTABLE; INJECTION

PROCAINAMIDE HYDROCHLORIDE

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	100MG/ML	A089029 001	Apr 17, 1986
	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 INC	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	
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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	
	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	

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PROCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

PROCAINE HYDROCHLORIDE

	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
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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
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PROCHLORPERAZINE

ALPHARMA US PHARMS	EQ 10MG BASE/ML	A087153 001	Jun 08, 1982
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PROCHLORPERAZINE EDISYLATE

MORTON GROVE	EQ 10MG BASE/ML	A088598 001	Oct 25, 1984
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INJECTABLE; INJECTION

COMPAZINE

+ GLAXOSMITHKLINE	EQ 5MG BASE/ML **	N010742 002
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PROCHLORPERAZINE

BAXTER HLTHCARE	EQ 5MG BASE/ML	A087759 001	Oct 01, 1982
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PROCHLORPERAZINE EDISYLATE

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
WEST-WARD PHARMS INT	EQ 5MG BASE/ML	A089523 001	May 03, 1988
WYETH AYERST	EQ 5MG BASE/ML	A086348 001	

SYRUP; ORAL

COMPAZINE

GLAXOSMITHKLINE	EQ 5MG BASE/5ML	N011188 001
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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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

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

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
SANDOZ	EQ 25MG BASE	A040101 003	Jul 19, 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 PHARMS	300MG	N019781 003	Oct 15, 1999
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INJECTABLE;INJECTION

PROGESTERONE

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	
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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

BAXTER HLTHCARE CORP	25MG/ML	N010349 008	
	50MG/ML	N010349 006	

SYRUP;ORAL

SPARINE

WYETH AYERST	10MG/5ML	N010942 003	
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TABLET;ORAL

SPARINE

WYETH AYERST	10MG	N010348 006	
	25MG	N010348 001	

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

6-354(of 430)

** See List Footnote

PROMAZINE HYDROCHLORIDETABLET; ORAL
SPARINE

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	
AKORN	25MG/ML	A083955 002	
	50MG/ML	A083955 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
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
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ZIPAN-50

ALTANA	50MG/ML	A083997 002
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SUPPOSITORY; RECTAL

PHENERGAN

+	MYLAN	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

SYRUP; ORAL

MYMETHAZINE FORTIS

USL PHARMA	25MG/5ML	A087996 001	Jan 18, 1983
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PROMETH FORTIS

ALPHARMA US PHARMS	25MG/5ML	A084772 001
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PROMETH PLAIN

ACTAVIS MID ATLANTIC	6.25MG/5ML	A085953 001
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PROMETHAZINE

CENCI	6.25MG/5ML	A089013 001	Sep 20, 1985
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PROMETHAZINE HYDROCHLORIDE

KV PHARM	6.25MG/5ML	A085388 001
	25MG/5ML	A085385 001
PHARM ASSOC	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	

DISCONTINUED DRUG PRODUCT LIST

6-355(of 430)

** See List Footnote

PROMETHAZINE HYDROCHLORIDE

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

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	

LANNETT

	12.5MG	A080949	001	
	25MG	A080949	002	
	50MG	A080949	003	

MYLAN

	12.5MG	A091054	001	Aug 30, 2011
	25MG	A091054	002	Aug 30, 2011
	50MG	A091054	003	Aug 30, 2011

PVT FORM

	12.5MG	A083214	001	
	25MG	A083658	001	

SANDOZ

	12.5MG	A084176	002	May 22, 2009
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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
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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	
	50MG	A083176	001	

PROPAFENONE HYDROCHLORIDE

TABLET; ORAL

PROPAFENONE HYDROCHLORIDE

NESHER PHARMS

	150MG	A076193	001	Feb 07, 2002
	225MG	A076193	002	Feb 07, 2002
	300MG	A076193	003	Feb 07, 2002

RYTHMOL

+ GLAXOSMITHKLINE LLC

	150MG	N019151	001	Nov 27, 1989
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+ 225MG

	N019151	003	Nov 20, 1992
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+ 300MG

	N019151	002	Nov 27, 1989
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PROPANTHELINE BROMIDE

INJECTABLE; INJECTION

PRO-BANTHINE

GD SEARLE LLC

	30MG/VIAL	N008843	001	
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TABLET; ORAL

PRO-BANTHINE

+ SHIRE

	7.5MG **	N008732	003	
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+ 15MG **

	N008732	002	
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PROPANTHELINE BROMIDE

ASCOT

	15MG	A087663	001	Oct 25, 1982
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Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

PROPANTHELINE BROMIDE

TABLET; ORAL

PROPANTHELINE BROMIDE

HEATHER	15MG	A085780	001	
HIKMA	7.5MG	A080927	001	
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	

PROPIOLACTONE

SOLUTION; IRRIGATION

BETAPRONE

FOREST LABS	N/A	N011657	001	
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PROPIOMAZINE HYDROCHLORIDE

INJECTABLE; INJECTION

LARGON

WEST-WARD PHARMS INT	20MG/ML	N012382	002	
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PROPOFOL

INJECTABLE; INJECTION

DIPRIVAN

FRESENIUS KABI USA	10MG/ML	N019627	001	Oct 02, 1989
PROPOFOL				
TEVA PARENTERAL	10MG/ML	A075392	001	Sep 19, 2000
WEST-WARD PHARMS INT	10MG/ML	A074848	001	Apr 19, 2005

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	

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PROPOXYPHENE HYDROCHLORIDE

CAPSULE; ORAL

PROPOXYPHENE HYDROCHLORIDE

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	
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TABLET; ORAL

DARVON-N

XANODYNE PHARM	100MG	N016862	002	
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PROPRANOLOL HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

PROPRANOLOL HYDROCHLORIDE

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
MYLAN	60MG	A078022	001	Feb 15, 2007
	80MG	A078022	002	Feb 15, 2007
	120MG	A078022	003	Feb 15, 2007
	160MG	A078022	004	Feb 15, 2007
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
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INJECTABLE; INJECTION

PROPRANOLOL HYDROCHLORIDE

+ BAXTER HLTHCARE CORP	1MG/ML	N016419	001	
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

MORTON GROVE	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
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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

ANI PHARMS INC	60MG	A071791	001	Jul 15, 1987
	90MG	A071977	001	Apr 06, 1988
DAVA PHARMS INC	10MG	A070125	001	Jul 30, 1985

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

PROPRANOLOL HYDROCHLORIDE

TABLET;ORAL

PROPRANOLOL HYDROCHLORIDE

	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
	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
PAR PHARM	90MG	A071288	001	Oct 22, 1986
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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

PROPRANOLOL HYDROCHLORIDE

TABLET; ORAL

PROPRANOLOL HYDROCHLORIDE

60MG	A070143 001	Jan 15, 1987	
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
YAOPHARMA CO LTD	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

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 INC	50MG	A080215 001
CHARTWELL RX	50MG	A084543 001
HALSEY	50MG	A080015 001
IMPAX LABS	50MG	A080159 001
LANNETT	50MG	A080016 001
LILLY	50MG	N006213 001
QUAGEN	50MG	A080154 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

+	LILLY	10MG/ML **	N006460 002	
	PHARMACIA AND UPJOHN	50MG/VIAL	N007413 001	
		250MG/VIAL	N007413 002	Aug 02, 1984
	WEST-WARD PHARMS INT	10MG/ML	A089474 001	Nov 05, 1986
		10MG/ML	A089475 001	Nov 05, 1986

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

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

PROTOKYLLOL HYDROCHLORIDETABLET;ORAL
VENTAIRE

SANOFI AVENTIS US 2MG A083459 001

PROTRIPTYLINE HYDROCHLORIDETABLET;ORAL
VIVACTILTEVA WOMENS 5MG ** N016012 001
10MG ** N016012 002PSEUDOEPHEDRINE HYDROCHLORIDECAPSULE, EXTENDED RELEASE;ORAL
NOVAFEDSANOFI AVENTIS US 120MG N017603 001
SUDAFED 12 HOUR
+ GLAXOSMITHKLINE 120MG ** N017941 002PSEUDOEPHEDRINE HYDROCHLORIDE; TRIPROLIDINE HYDROCHLORIDECAPSULE, EXTENDED RELEASE;ORAL
ACTIFEDGLAXOSMITHKLINE 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
FOSUN PHARMA 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, 1991PSEUDOEPHEDRINE POLISTIREXSUSPENSION, EXTENDED RELEASE;ORAL
PSEUDO-12

UCB INC EQ 60MG HYDROCHLORIDE/5ML N019401 001 Jun 19, 1987

PSEUDOEPHEDRINE SULFATETABLET, EXTENDED RELEASE;ORAL
AFRINOL

+ SCHERING PLOUGH 120MG N018191 001

PYRAZINAMIDE

TABLET;ORAL

PYRAZINAMIDE
DAVA PHARMS INC 500MG A080157 001

DISCONTINUED DRUG PRODUCT LIST

6-361(of 430)

** See List Footnote

PYRIDOSTIGMINE BROMIDE

TABLET; ORAL

PYRIDOSTIGMINE BROMIDE

ANI PHARMS INC	30MG	A040512 002	Jul 20, 2005
	60MG	A040512 001	Oct 08, 2003
IMPAX LABS INC	60MG	A040457 001	Dec 26, 2002
SOLVAY	30MG	A089572 001	Nov 27, 1990
US ARMY	30MG	N020414 001	Feb 05, 2003

PYRIDOXINE HYDROCHLORIDE

INJECTABLE; INJECTION

HEXA-BETALIN

LILLY	100MG/ML	A080854 001	
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PYRIDOXINE HYDROCHLORIDE

AKORN	100MG/ML	A087967 001	Oct 01, 1982
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	
LUITPOLD	100MG/ML	A080669 001	
MYLAN INSTITUTIONAL	100MG/ML	A204879 001	Jul 14, 2016
WATSON LABS	100MG/ML	A083760 001	

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	
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PYRITHIONE ZINC

LOTION; TOPICAL

HEAD & SHOULDERS CONDITIONER

WARNER CHILCOTT	0.3%	N019412 002	Mar 10, 1986
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PYRVINIUM PAMOATE

SUSPENSION; ORAL

POVAN

PARKE DAVIS	EQ 50MG BASE/5ML	N011964 001	
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TABLET; ORAL

POVAN

PARKE DAVIS	EQ 50MG BASE	N012485 002	
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QUAZEPAM

TABLET; ORAL

DORAL

GALT PHARMS	7.5MG	N018708 003	Feb 26, 1987
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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
MYLAN PHARMS INC	EQ 25MG BASE	A090323 001	Mar 27, 2012

SEROQUEL

+ ASTRAZENECA PHARMS

EQ 150MG BASE **	N020639 004	Dec 20, 1998
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TABLET, EXTENDED RELEASE; ORAL

QUETIAPINE FUMARATE

AMNEAL PHARMS	EQ 400MG BASE	A211405 001	Oct 26, 2018
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DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

QUINAPRIL HYDROCHLORIDE

TABLET; ORAL

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 INC	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
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
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
YAOPHARMA CO LTD	EQ 5MG BASE	A076803 001	Mar 02, 2005
	EQ 10MG BASE	A076803 002	Mar 02, 2005
	EQ 20MG BASE	A076803 003	Mar 02, 2005
	EQ 40MG BASE	A076803 004	Mar 02, 2005

QUINESTROL

TABLET; ORAL

ESTROVIS

PARKE DAVIS	0.1MG	N016768 002
	0.2MG	N016768 003

QUINETHAZONE

TABLET; ORAL

HYDROMOX

LEDERLE	50MG	N013264 001
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QUINETHAZONE; RESERPINE

TABLET; ORAL

HYDROMOX R

LEDERLE	50MG; 0.125MG	N013927 001
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QUINIDINE GLUCONATE

INJECTABLE; INJECTION

QUINIDINE GLUCONATE

+ LILLY

80MG/ML	N007529 002	Feb 10, 1989
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TABLET; ORAL

QUINACT

BAYER HLTHCARE	266MG	A085978 001
	400MG	A086099 001

TABLET, EXTENDED RELEASE; ORAL

DURAQUIN

WARNER CHILCOTT	330MG	N017917 001
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QUINAGLUTE

+ BAYER HLTHCARE

324MG	N016647 001
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QUINALAN

LANNETT

324MG	A088081 001	Feb 10, 1986
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QUINATIME

WATSON LABS

324MG	A087448 001
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QUINIDINE GLUCONATE

ANI PHARMS INC

324MG	A087810 001	Sep 29, 1982
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Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

QUINIDINE GLUCONATE

TABLET, EXTENDED RELEASE;ORAL

QUINIDINE GLUCONATE

ASCOT	324MG	A088582 001	Jun 17, 1985
AUROLIFE PHARMA LLC	324MG	A089894 001	Dec 15, 1988
CYCLE PHARMS LTD	324MG	A088431 001	Jan 06, 1984
HALSEY	324MG	A089476 001	Apr 10, 1987
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	
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QUINIDINE SULFATE

CAPSULE;ORAL

CIN-QUIN

SOLVAY	200MG	A085296 001	
	300MG	A085297 001	

QUINIDINE SULFATE

LILLY	200MG	A085103 001	
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TABLET;ORAL

CIN-QUIN

SOLVAY	100MG	A085299 001	
	200MG	A084932 001	
	300MG	A085298 001	

QUINIDINE SULFATE

BARR	200MG	A084177 001	
CONTRACT PHARMACAL	200MG	A083808 001	
CYCLE PHARMS LTD	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 PHARMS	200MG	A083862 001	
IMPAX LABS	200MG	A083347 001	
IVAX SUB TEVA PHARMS	200MG	A084549 001	
KING PHARMS	200MG	A085175 001	
KV PHARM	200MG	A085276 001	
LANNETT	200MG	A083743 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	
	300MG	A089839 001	Sep 29, 1988
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	A085140 002	
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	
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DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

QUINIDINE SULFATE

TABLET, EXTENDED RELEASE;ORAL

QUINIDINE SULFATE

ACP NIMBLE

300MG

A040045 001 Jun 30, 1994

QUININE SULFATE

CAPSULE;ORAL

QUININE SULFATE

MYLAN PHARMS INC

324MG

A202581 001 Dec 14, 2012

RABEPRAZOLE SODIUM

TABLET, DELAYED RELEASE;ORAL

ACIPHEX

+ EISAI INC

10MG **

N020973 001 May 29, 2002

RABEPRAZOLE SODIUM

MYLAN

20MG

A076885 001 Nov 08, 2013

RAMIPRIL

CAPSULE;ORAL

RAMIPRIL

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

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

WATSON LABS

5MG

A076549 003 Oct 24, 2005

YAOPHARMA CO LTD

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

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

MYLAN PHARMS INC

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

MYLAN

EQ 150MG BASE

A075564 001 Oct 27, 2000

EQ 300MG BASE

A075564 002 Oct 27, 2000

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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

RANITIDINE HYDROCHLORIDE

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

ZANTAC IN PLASTIC CONTAINER

TELIGENT 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

APOTEX INC EQ 15MG BASE/ML A077602 001 Sep 17, 2007

RANBAXY EQ 15MG BASE/ML A078448 001 Dec 13, 2007

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 INC 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

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

MYLAN EQ 75MG BASE A075497 001 Jan 14, 2000

EQ 150MG BASE A074023 001 Aug 22, 1997

EQ 150MG BASE A074552 001 Jul 30, 1998

EQ 300MG BASE A074023 002 Aug 22, 1997

EQ 300MG BASE A074552 002 Jul 30, 1998

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 150MG BASE A210010 001 Aug 01, 2018

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

WATSON LABS EQ 150MG BASE A074864 001 Oct 20, 1997

EQ 300MG BASE A074864 002 Oct 20, 1997

WOCKHARDT 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

ZANTAC 150

+ GLAXO GRP LTD EQ 150MG BASE ** N018703 001 Jun 09, 1983

ZANTAC 300

+ GLAXO GRP LTD EQ 300MG BASE ** N018703 002 Dec 09, 1985

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

+ SANOFI US EQ 75MG BASE ** N020745 001 Feb 26, 1998

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

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

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

FERNDALE 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

MYLAN

0.5MG

A090252 001 Aug 23, 2013

1MG

A090252 002 Jan 22, 2014

2MG

A090252 003 Jan 22, 2014

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

RESCINNAMINE

CAPSULE; ORAL

CINNASIL

PANRAY

0.5MG

A084736 001

TABLET; ORAL

MODERIL

PFIZER

0.25MG

N010686 003

0.5MG

N010686 006

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 PHARMS LTD

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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

RESERPINE

TABLET;ORAL

RESERPINE

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
	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

RIBAVIRIN

CAPSULE;ORAL

REBETOL

MERCK SHARP DOHME	200MG**Indicated for use and comarketed with Interferon ALFA-2B, Recombinant (INTRON A), as Rebetrone Combination Therapy**	N020903 001	Jun 03, 1998
RIBASPHERE			
KADMON PHARMS LLC	200MG	A076203 001	Apr 06, 2004
RIBAVIRIN			
CASI PHARMS INC	200MG	A076192 001	Apr 06, 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
HERITAGE PHARMA	200MG	A077053 001	Dec 05, 2005
KADMON PHARMS LLC	200MG	A077456 001	Dec 05, 2005
	400MG	A077456 002	Dec 05, 2005
	600MG	A077456 003	Dec 05, 2005
ZYDUS PHARMS USA	400MG	A077094 002	Mar 16, 2007

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

RIBAVIRINTABLET;ORAL
RIBAVIRIN

500MG	A077094	004	Apr 18, 2008
600MG	A077094	003	Mar 16, 2007

RILUZOLETABLET;ORAL
RILUZOLE

DAITO PHARMS CO LTD	50MG	A204430	001	Oct 16, 2018
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RIMANTADINE HYDROCHLORIDESYRUP;ORAL
FLUMADINE

FOREST LABS	50MG/5ML	N019650	001	Sep 17, 1993
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TABLET;ORAL

RIMANTADINE HYDROCHLORIDE

HERITAGE PHARMA	100MG	A076375	001	Jan 14, 2003
IMPAX LABS INC	100MG	A075916	001	Nov 02, 2001

RIMEXOLONESUSPENSION/DROPS;OPHTHALMIC
VEXOL

EYEVANCE	1%	N020474	001	Dec 30, 1994
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RISEDRONATE SODIUM

TABLET;ORAL

ACTONEL

+ APIL 75MG **

N020835	004	Apr 16, 2007
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RISEDRONATE SODIUM

HANGZHOU BINJIANG	35MG	A207516	001	Feb 15, 2019
MYLAN	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

RISPERIDONE

SOLUTION;ORAL

RISPERIDONE

ANI PHARMS INC	1MG/ML	A076440	001	Jan 30, 2009
LANNETT CO INC	1MG/ML	A202386	001	Jan 12, 2015
PRECISION DOSE	1MG/ML	A076797	001	Jun 28, 2010
TORRENT	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
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RISPERIDONE

HERITAGE PHARMA	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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

RISPERIDONETABLET; ORAL
RISPERIDONE

	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
	3MG	A078740 005	May 29, 2009
	4MG	A078740 006	May 29, 2009
TABLET, ORALLY DISINTEGRATING; ORAL			
RISPERIDONE			
MYLAN PHARMS INC	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	
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RITONAVIR

CAPSULE; ORAL

NORVIR

ABBOTT	100MG	N020680 001	Mar 01, 1996
+ ABBVIE	100MG **	N020945 001	Jun 29, 1999

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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

RIVASTIGMINE TARTRATE

SOLUTION; ORAL

EXELON

NOVARTIS

EQ 2MG BASE/ML

N021025 001 Apr 21, 2000

RIZATRIPTAN BENZOATE

TABLET; ORAL

MAXALT

+ MERCK

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

TABLET, ORALLY DISINTEGRATING; ORAL

MAXALT-MLT

+ MERCK

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

MYLAN PHARMS INC

EQ 5MG BASE

A078173 001 Dec 31, 2012

EQ 10MG BASE

A078173 002 Dec 31, 2012

ROCURONIUM BROMIDE

INJECTABLE; INJECTION

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

ROFLUMILAST

TABLET; ORAL

ROFLUMILAST

MYLAN

500MCG

A208257 001 Jul 13, 2018

ROLAPITANT HYDROCHLORIDE

EMULSION; INTRAVENOUS

VARUBI

+ TERSERA THERAPS LLC

EQ 166.5MG BASE/92.5ML (EQ 1.8MG
BASE/ML)

N208399 001 Oct 25, 2017

ROPINIROLE HYDROCHLORIDE

TABLET; ORAL

ROPINIROLE HYDROCHLORIDE

ACP NIMBLE

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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

ROPINIROLE HYDROCHLORIDE

TABLET;ORAL

ROPINIROLE HYDROCHLORIDE

EQ 4MG BASE

A077852 006 May 05, 2008

EQ 5MG BASE

A077852 007 May 19, 2008

TABLET, EXTENDED RELEASE;ORAL

REQUIP XL

+ GLAXOSMITHKLINE LLC EQ 3MG BASE **

N022008 002 Jun 13, 2008

+ EQ 4MG BASE

N022008 003 Jun 13, 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

EQ 8MG BASE

A200462 005 Oct 15, 2012

EQ 12MG BASE

A200462 006 Oct 15, 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

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

+ SB PHARMCO

EQ 8MG BASE

N021071 004 May 25, 1999

ROSIGLITAZONE MALEATE

ANI PHARMS INC

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

TABLET;ORAL

ROSUVASTATIN CALCIUM

AMNEAL PHARMS CO

5MG

A208850 001 Oct 16, 2018

10MG

A208850 002 Oct 16, 2018

20MG

A208850 003 Oct 16, 2018

40MG

A208850 004 Oct 16, 2018

MYLAN

5MG

A079161 001 Jul 19, 2016

10MG

A079161 002 Jul 19, 2016

20MG

A079161 003 Jul 19, 2016

40MG

A079161 004 Jul 19, 2016

SCIEGEN PHARMS INC

5MG

A206381 001 Apr 24, 2019

10MG

A206381 002 Apr 24, 2019

20MG

A206381 003 Apr 24, 2019

40MG

A206381 004 Apr 24, 2019

TEVA PHARMS USA

5MG

A079166 001 Jul 19, 2016

10MG

A079166 002 Jul 19, 2016

20MG

A079166 003 Jul 19, 2016

40MG

A079166 004 Jul 19, 2016

UMEDICA LABS PVT LTD

5MG

A207626 001 Apr 09, 2019

10MG

A207626 002 Apr 09, 2019

20MG

A207626 003 Apr 09, 2019

40MG

A207626 004 Apr 09, 2019

ZYDUS PHARMS

5MG

A206513 001 Mar 01, 2019

10MG

A206513 002 Mar 01, 2019

20MG

A206513 003 Mar 01, 2019

40MG

A206513 004 Mar 01, 2019

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

RUFINAMIDE

TABLET; ORAL

BANZEL

+ EISAI INC

100MG **

N021911 001 Nov 14, 2008

RUFINAMIDE

MYLAN

200MG

A205095 001 May 16, 2016

400MG

A205095 002 May 16, 2016

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

SALMETEROL XINAFOATE

AEROSOL, METERED; INHALATION

SEREVENT

GLAXOSMITHKLINE

EQ 0.021MG BASE/INH

N020236 001 Feb 04, 1994

SAPROPTERIN DIHYDROCHLORIDE

POWDER; ORAL

SAPROPTERIN DIHYDROCHLORIDE

PAR PHARM INC

500MG/PACKET

A210027 001 Aug 20, 2019

TABLET; ORAL

SAPROPTERIN DIHYDROCHLORIDE

PAR PHARM INC

100MG

A207200 001 May 10, 2019

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

SARALASIN ACETATE

INJECTABLE; INJECTION

SARENIN

PROCTER AND GAMBLE

EQ 0.6MG BASE/ML

N018009 001

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

WHITEWORTH TOWN PLSN

100MG

A085798 001

WYETH AYERST

100MG

A086390 001

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

SECOBARBITAL SODIUM

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

+ SOMERSET 5MG N020647 001 May 15, 1996

SELEGILINE HYDROCHLORIDE

DAVA PHARMS INC 5MG A075352 001 Nov 30, 1998

LANNETT CO INC 5MG A075145 001 Sep 15, 2003

TABLET; ORAL

SELEGILINE HYDROCHLORIDE

ACP NIMBLE 5MG A074744 001 Jan 27, 1997

5MG A074756 001 Nov 25, 1998

ALLIED 5MG A074672 001 Apr 01, 1997

CHARTWELL MOLECULES 5MG A074565 001 Aug 02, 1996

5MG A074641 001 Aug 02, 1996

G AND W LABS INC 5MG A074537 001 Aug 02, 1996

MYLAN 5MG A074866 001 Nov 26, 1997

+ SOMERSET 5MG ** N019334 001 Jun 05, 1989

SELENIUM SULFIDE

LOTION; SHAMPOO; TOPICAL

EXSEL

ALLERGAN HERBERT 2.5% A083892 001

SELENIUM SULFIDE

ACP NIMBLE 2.5% A086209 001

ACTAVIS MID ATLANTIC 2.5% A084394 001

IVAX PHARMS 2.5% A085777 001

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

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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

SERTRALINE HYDROCHLORIDE

CONCENTRATE; ORAL

SERTRALINE HYDROCHLORIDE

ACI HEALTHCARE LTD	EQ 20MG BASE/ML	A076934	001	Jun 30, 2006
RANBAXY LABS LTD	EQ 20MG BASE/ML	A078053	001	Feb 05, 2007

TABLET; ORAL

SERTRALINE HYDROCHLORIDE

ACI HEALTHCARE LTD	EQ 25MG BASE	A076881	001	Feb 06, 2007
	EQ 50MG BASE	A076881	002	Feb 06, 2007
	EQ 100MG BASE	A076881	003	Feb 06, 2007
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
CADILA	EQ 25MG BASE	A077106	001	Feb 06, 2007
	EQ 50MG BASE	A077106	002	Feb 06, 2007
	EQ 100MG BASE	A077106	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
FOSUN PHARMA	EQ 25MG BASE	A077713	001	Feb 06, 2007
	EQ 50MG BASE	A077713	002	Feb 06, 2007
	EQ 100MG BASE	A077713	003	Feb 06, 2007
HERITAGE PHARMA	EQ 25MG BASE	A076465	001	Aug 11, 2006
	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	A076465	002	Aug 11, 2006
	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	A076465	003	Aug 11, 2006
	EQ 100MG BASE	A077299	003	Feb 06, 2007
	EQ 100MG BASE	A077345	003	Feb 06, 2007
	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	EQ 25MG BASE	A076671	001	Feb 06, 2007
	EQ 50MG BASE	A076671	002	Feb 06, 2007
	EQ 100MG BASE	A076671	003	Feb 06, 2007
MYLAN PHARMS INC	EQ 25MG BASE	A076540	001	Mar 20, 2007
	EQ 25MG BASE	A078626	001	Jan 31, 2008
	EQ 50MG BASE	A076540	002	Mar 20, 2007
	EQ 50MG BASE	A078626	002	Jan 31, 2008
	EQ 100MG BASE	A076540	003	Mar 20, 2007
	EQ 100MG BASE	A078626	003	Jan 31, 2008
SCIEGEN PHARMS INC	EQ 25MG BASE	A076442	001	Apr 30, 2007
	EQ 50MG BASE	A076442	002	Apr 30, 2007
	EQ 100MG BASE	A076442	003	Apr 30, 2007
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

ZOLOFT

+	PFIZER	EQ 150MG BASE **	N019839	003	Dec 30, 1991
+		EQ 200MG BASE **	N019839	004	Dec 30, 1991

SEVELAMER HYDROCHLORIDE

CAPSULE; ORAL

RENAGEL

GENZYME	403MG	N020926	001	Oct 30, 1998
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DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

SIBUTRAMINE HYDROCHLORIDE

CAPSULE; ORAL

MERIDIA

ABBOTT	5MG	N020632 001	Nov 22, 1997
	10MG	N020632 002	Nov 22, 1997
	15MG	N020632 003	Nov 22, 1997

SILDENAFIL CITRATE

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

SILVER SULFADIAZINE

CREAM; TOPICAL

SSD AF

DR REDDYS LA	1%	N018578 003	Jul 11, 1990
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DRESSING; TOPICAL

SILDAFLO

FRANKLIN PHARMS	1%	N019608 001	Nov 30, 1989
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SIMEPREVIR SODIUM

CAPSULE; ORAL

OLYSIO

+ JANSSEN PRODS	EQ 150MG BASE	N205123 001	Nov 22, 2013
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SIMETHICONE-CELLULOSE

SUSPENSION; ORAL

SONORX

BRACCO	7.5MG/ML	N020773 001	Oct 29, 1998
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SIMVASTATIN

TABLET; ORAL

SIMVASTATIN

HISUN PHARM HANGZHOU	10MG	A206557 001	Nov 13, 2017
	20MG	A206557 002	Nov 13, 2017
	40MG	A206557 003	Nov 13, 2017
	80MG	A206557 004	Nov 13, 2017
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
MYLAN PHARMS INC	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
YAOPHARMA CO LTD	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
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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

SIMVASTATIN; SITAGLIPTIN PHOSPHATE

TABLET; ORAL

JUVISYNC

+	40MG;EQ 50MG BASE **	N202343 006	Sep 18, 2012
+	40MG;EQ 100MG BASE **	N202343 003	Oct 07, 2011

SIROLIMUS

TABLET; ORAL

RAPAMUNE

+	PF PRISM CV	5MG **	N021110 003	Feb 23, 2004
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SODIUM BENZOATE; SODIUM PHENYLACETATE

SOLUTION; ORAL

UCEPHAN

B BRAUN	100MG/ML;100MG/ML	N019530 001	Dec 23, 1987
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SODIUM BICARBONATE

INJECTABLE; INJECTION

SODIUM BICARBONATE

HOSPIRA INC	0.5MEQ/ML	A202679 001	Mar 07, 2017
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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
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SODIUM CHLORIDE

INJECTABLE; INJECTION

BACTERIOSTATIC SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

ABRAXIS PHARM	9MG/ML	A088909 001	Feb 07, 1985
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SODIUM CHLORIDE

ABBOTT	20GM/100ML	N017013 001
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B BRAUN	20GM/100ML	N017038 001
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SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER

B BRAUN	450MG/100ML	N018184 001
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MILES	450MG/100ML	N018503 001
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SODIUM CHLORIDE 0.9%

+	MEDEFIL INC	18MG/2ML (9MG/ML)	N202832 002	Jan 06, 2012
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+		22.5MG/2.5ML (9MG/ML)	N202832 003	Jan 06, 2012
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+		27MG/3ML (9MG/ML)	N202832 004	Jan 06, 2012
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+		45MG/5ML (9MG/ML)	N202832 005	Jan 06, 2012
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SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

ABBOTT	9MG/ML	N019218 001	Jul 13, 1984
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+	ICU MEDICAL INC	9MG/ML	N019217 001	Jul 13, 1984
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+	LIEBEL-FLARSHEIM	405MG/50ML (9MG/ML)	N021569 001	Jul 27, 2006
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+	MEDEFIL INC	9MG/ML (9MG/ML)	N202832 001	Jan 06, 2012
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MILES	900MG/100ML	N018502 001
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SODIUM CHLORIDE 23.4% IN PLASTIC CONTAINER

+	ABRAXIS PHARM	234MG/ML **	N019329 001	Apr 22, 1987
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SOLUTION; IRRIGATION

SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER

BAXTER HLTHCARE	450MG/100ML	N017864 001
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	450MG/100ML	N018497 001	Feb 19, 1982
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HOSPIRA	450MG/100ML	N017670 001
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	450MG/100ML	N018380 001
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SODIUM CHLORIDE IN PLASTIC CONTAINER

MILES	900MG/100ML	N018247 001
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SODIUM CHROMATE CR-51

INJECTABLE; INJECTION

CHROMITOPE SODIUM

BRACCO	2mCi/VIAL	N013993 002
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	200uCi/ML	N013993 001
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SODIUM CHROMATE CR 51

CURIUM	100uCi/ML	N016708 001
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DISCONTINUED DRUG PRODUCT LIST

6-378(of 430)

** See List Footnote

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			
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
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SODIUM IODIDE I-123

CAPSULE; ORAL

SODIUM IODIDE I 123

CARDINAL HEALTH 418	400uCi	N018671 003	May 27, 1982
GE HEALTHCARE	100uCi	N017630 001	

SOLUTION; ORAL

SODIUM IODIDE I 123

GE HEALTHCARE	2mCi/ML **	N017630 002	
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SODIUM IODIDE I-131

CAPSULE; ORAL

IODOTOPE

BRACCO	1-130mCi	N010929 001	
	1-150mCi	N010929 003	

SODIUM IODIDE I 131

CIS	50uCi	N017316 001	
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	100uCi	N017316 002	
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CURIUM	0.8-100mCi	N016515 002	
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+ CURIUM	0.8-100mCi	N016517 001	
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	15-100uCi	N016517 002	
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JUBILANT DRAXIMAGE	2-200mCi	N021305 004	Nov 18, 2004
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SOLUTION; ORAL

HICON

JUBILANT DRAXIMAGE	1-250mCi/0.25ML	N021305 002	Jan 24, 2003
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	1-500mCi/0.5ML	N021305 003	Jan 24, 2003
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	1-1000mCi/ML	N021305 005	Apr 04, 2006
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IODOTOPE

BRACCO	7-106mCi/BOT	N010929 002	
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SODIUM IODIDE I 131

CIS	50mCi/ML	N017315 001	
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+ CURIUM	3.5-150mCi/VIAL	N016515 001	
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SODIUM LACTATE

INJECTABLE; INJECTION

SODIUM LACTATE 0.167 MOLAR IN PLASTIC CONTAINER

B BRAUN	1.87GM/100ML	N018186 001	
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BAXTER HLTHCARE	1.87GM/100ML	N016692 001	
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HOSPIRA	1.87GM/100ML	N018249 001	
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SODIUM LACTATE 1/6 MOLAR IN PLASTIC CONTAINER

B BRAUN	1.87GM/100ML	N020004 001	Apr 21, 1992
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SODIUM MONOFLUOROPHOSPHATE

GEL; DENTAL

EXTRA-STRENGTH AIM

CHESEBROUGH PONDS	1.2%	N019518 002	Aug 06, 1986
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PASTE; DENTAL

EXTRA-STRENGTH AIM

CHESEBROUGH PONDS	1.2%	N019518 001	Jun 03, 1987
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SODIUM NITROPRUSSIDE

INJECTABLE; INJECTION

NIPRIDE

ROCHE	50MG/VIAL	N017546 001	
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NITROPRESS

ABBOTT	50MG/VIAL	A071555 001	Nov 16, 1987
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+ ABBVIE	50MG/VIAL **	N018450 001	
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HOSPIRA	50MG/VIAL	A070566 001	Jun 09, 1986
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Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

SODIUM NITROPRUSSIDE

INJECTABLE; INJECTION

SODIUM NITROPRUSSIDE

ABRAXIS PHARM	50MG/VIAL	A070031 001	Jan 17, 1985
+ BAXTER HLTHCARE	50MG/VIAL **	N018581 001	Jul 28, 1982
SUN PHARM	25MG/ML	A210467 001	Nov 26, 2018
TEVA PARENTERAL	25MG/ML	A073465 001	Mar 30, 1992
VIRTUS PHARM	25MG/ML	A209834 001	Jun 26, 2018

SOLUTION; INTRAVENOUS

NIPRIDE RTU IN SODIUM CHLORIDE 0.9%

+ EXELA PHARMA SCS LLC	10MG/50ML (0.2MG/ML)	N209387 002	Dec 07, 2017
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SODIUM PHENYLBUTYRATE

TABLET; ORAL

SODIUM PHENYLBUTYRATE

ALVOGEN	500MG	A090910 001	Nov 18, 2011
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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

VISICOL

SALIX PHARMS	0.398GM;1.102GM	N021097 001	Sep 21, 2000
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SODIUM 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
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SODIUM POLYSTYRENE SULFONATE

POWDER; ORAL, RECTAL

KAYEXALATE

+ CONCORDIA	453.6GM/BOT **	N011287 001	
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SODIUM POLYSTYRENE SULFONATE

CITRUSPHRMA	454GM/BOT	A040909 001	Dec 03, 2008
WOCKHARDT	453.6GM/BOT	A088786 001	Sep 11, 1984

SUSPENSION; ORAL, RECTAL

SODIUM POLYSTYRENE SULFONATE

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	
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SODIUM TETRADECYL SULFATE

INJECTABLE; INJECTION

SOTRADECOL

+ ELKINS SINN	1% **	N005970 004	
+ ELKINS SINN	3% **	N005970 005	

SODIUM THIOSULFATE

INJECTABLE; INJECTION

SODIUM THIOSULFATE

US ARMY	250MG/ML	N020166 001	Feb 14, 1992
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SOMATREM

INJECTABLE; INJECTION

PROTROPIN

GENENTECH	5MG/VIAL	N019107 001	Oct 17, 1985
	10MG/VIAL	N019107 002	Oct 24, 1989

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

SOMATROPIN

INJECTABLE; INJECTION

ACCRETROPIN					
EMERGENT	5MG/ML (5MG/ML)		N021538	001	Jan 23, 2008
ASELLACRIN 10					
SERONO	10 IU/VIAL		N017726	001	
ASELLACRIN 2					
SERONO	2 IU/VIAL		N017726	002	Jul 21, 1983
BIO-TROPIN					
FERRING	4.8MG/VIAL		N019774	001	May 25, 1995
CRESCORMON					
GENENTECH	4 IU/VIAL		N017992	001	
GENOTROPIN PRESERVATIVE FREE					
+ PHARMACIA	1.5MG/VIAL		N020280	004	Aug 24, 1995
HUMATROPE					
LILLY	2MG/VIAL		N019640	001	Jun 23, 1987
NORDITROPIN					
NOVO NORDISK INC	5MG/1.5ML		N021148	001	Jun 20, 2000
	10MG/1.5ML		N021148	002	Jun 20, 2000
	15MG/1.5ML		N021148	003	Jun 20, 2000
NORDITROPIN NORDIFLEX					
NOVO NORDISK INC	5MG/1.5ML		N021148	004	Oct 01, 2004
	10MG/1.5ML		N021148	005	Oct 01, 2004
	15MG/1.5ML		N021148	006	Oct 01, 2004
	30MG/3ML		N021148	007	Mar 10, 2009
NUTROPIN					
GENENTECH	5MG/VIAL		N020168	001	Nov 17, 1993
	10MG/VIAL		N020168	002	Nov 17, 1993
NUTROPIN AQ					
GENENTECH	10MG/2ML (5MG/ML)		N020522	001	Dec 29, 1995
NUTROPIN AQ PEN					
+ GENENTECH	10MG/2ML (5MG/ML)		N020522	002	Apr 22, 2002
+ GENENTECH	20MG/2ML (10MG/ML)		N020522	006	Jan 03, 2008
SAIZEN					
EMD SERONO	4MG/VIAL		N019764	005	Jan 16, 2007
	6MG/VIAL		N019764	001	Oct 08, 1996
SEROSTIM					
EMD SERONO	8.8MG/VIAL		N020604	004	Sep 06, 2001
INJECTABLE; SUBCUTANEOUS					
SEROSTIM LQ					
EMD SERONO	6MG/0.5ML (6MG/0.5ML)		N020604	005	Feb 11, 2005

SOMATROPIN RECOMBINANT

INJECTABLE; INJECTION

NORDITROPIN					
NOVO NORDISK INC	4MG/VIAL		N019721	001	May 08, 1995
	8MG/VIAL		N019721	002	May 08, 1995
NUTROPIN DEPOT					
GENENTECH	13.5MG/VIAL		N021075	001	Dec 22, 1999
	18MG/VIAL		N021075	002	Dec 22, 1999
	22.5MG/VIAL		N021075	003	Dec 22, 1999
VALTROPIN					
LG CHEM LTD	5MG/VIAL		N021905	001	Apr 19, 2007
ZORBTIVE					
EMD SERONO	4MG/VIAL		N021597	001	Dec 01, 2003
	5MG/VIAL		N021597	002	Dec 01, 2003
	6MG/VIAL		N021597	003	Dec 01, 2003

SORBITOL

SOLUTION; IRRIGATION

SORBITOL 3% IN PLASTIC CONTAINER					
BAXTER HLTHCARE	3GM/100ML		N018512	001	May 27, 1982

SOTALOL HYDROCHLORIDE

TABLET; ORAL

BETAPACE					
COVIS PHARMA BV	320MG		N019865	004	Oct 30, 1992
BETAPACE AF					
COVIS PHARMA BV	40MG		N021151	006	Apr 02, 2003
	60MG		N021151	007	Apr 02, 2003
	100MG		N021151	005	Mar 14, 2003

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

SOTALOL HYDROCHLORIDE

TABLET; ORAL

SOTALOL HYDROCHLORIDE

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
MYLAN	80MG	A075237 001	May 01, 2000
	80MG	A075725 001	Dec 19, 2000
	120MG	A075237 002	May 01, 2000
	120MG	A075725 002	Dec 19, 2000
	160MG	A075237 003	May 01, 2000
	160MG	A075725 003	Dec 19, 2000
	240MG	A075237 004	May 01, 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

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
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
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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

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

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
AUROBINDO PHARMA LTD	25MG	A202187 001	Mar 06, 2014
	50MG	A202187 002	Mar 06, 2014
	100MG	A202187 003	Mar 06, 2014
CASI PHARMS INC	25MG	A086809 001	
IVAX PHARMS	25MG	A087108 001	
LEDERLE	25MG	A087634 001	
MUTUAL PHARM	25MG	A087265 001	
MYLAN	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

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
MYLAN	15MG	A079069 001	Dec 29, 2008
	20MG	A079069 002	Dec 29, 2008
	30MG	A079069 003	Dec 29, 2008
	40MG	A079069 004	Dec 29, 2008
MYLAN LABS LTD	30MG	A078775 001	Jan 05, 2009
	40MG	A078775 002	Jan 05, 2009
ZERIT			
BRISTOL MYERS SQUIBB	5MG	N020412 001	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-MYERS SQUIBB	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			
+ HOSPIRA	100% (1ML)	N018801 001	Oct 27, 1982
WEST-WARD PHARMS INT	100% (20ML)	A206369 002	Sep 02, 2015
STERILE WATER FOR INJECTION IN PLASTIC CONTAINER			
B BRAUN	100%	N019077 001	Mar 02, 1984

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

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 INC 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

INTL MEDICATION 100MG/VIAL A085400 001 Feb 04, 1982

ORGANON USA INC 20MG/ML A080997 001

SUCOSTRIN

APOTHECON 20MG/ML N008847 001

100MG/ML N008847 003

SUCRALFATE

TABLET; ORAL

SUCRALFATE

NOSTRUM LABS INC 1GM A074415 001 Jun 08, 1998

SUFENTANIL CITRATE

INJECTABLE; INJECTION

SUFENTANIL CITRATE

WATSON LABS EQ 0.05MG BASE/ML A074406 001 Dec 15, 1995

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-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

AKORN 10% A083021 001

15% A083021 002

30% A083021 003

SOLA BARNES HIND 10% A084143 001

10% A084145 001

30% A084146 001

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

SULFACETAMIDE SODIUMSOLUTION/DROPS;OPHTHALMIC
SODIUM SULFACETAMIDE

	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			
AKORN	30%	A040216 001	May 25, 1999
ALCON PHARMS LTD	30%	A089068 001	May 05, 1987
PHARMAFAIR	10%	A088947 001	May 17, 1985
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

SULFACYTINETABLET;ORAL
RENOQUID

GLENWOOD	250MG	N017569 001	
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SULFADIAZINE

TABLET;ORAL

SULFADIAZINE

ABBVIE	300MG	N004125 005	
EVERYLIFE	500MG	A080088 001	
IMPAX LABS	500MG	A080081 001	
LANNETT	500MG	A080084 001	
LEDERLE	500MG	N004054 001	
+ LILLY	500MG	N004122 002	

SULFADIAZINE SODIUM

INJECTABLE; INJECTION

SULFADIAZINE SODIUM

LEDERLE	250MG/ML	N004054 002	
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SULFADIAZINE; SULFAMERAZINE

SUSPENSION;ORAL

SULFONAMIDES DUPLEX

LILLY	250MG/5ML;250MG/5ML	N006317 007	
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SULFAMETER

TABLET;ORAL

SULLA

BAYER HLTHCARE	500MG	N016000 002	
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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	
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TABLET;ORAL

GANTANOL

ROCHE	500MG	N012715 002	
GANTANOL-DS			
ROCHE	1GM	N012715 003	

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

6-385(of 430)

** See List Footnote

SULFAMETHOXAZOLE

TABLET; ORAL

SULFAMETHOXAZOLE

ASCOT	500MG	A087662	001	Oct 20, 1982
AUROLIFE PHARMA LLC	500MG	A085844	001	
BARR	500MG	A087189	001	Jul 25, 1983
HEATHER	500MG	A086163	001	
WATSON LABS	500MG	A085053	001	
	1GM	A086000	001	

UROBAK

SHIONOGI	500MG	A087307	001	
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SULFAMETHOXAZOLE; TRIMETHOPRIM

INJECTABLE; INJECTION

BACTRIM

+ SUN PHARM INDS INC	80MG/ML;16MG/ML **	N018374	001	
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SEPTRA

MONARCH PHARMS	80MG/ML;16MG/ML **	N018452	001	
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SULFAMETHOXAZOLE AND TRIMETHOPRIM

ABRAXIS PHARM	80MG/ML;16MG/ML	A070223	001	Dec 29, 1987
BEDFORD	80MG/ML;16MG/ML	A072383	001	Apr 29, 1992
HOSPIRA	80MG/ML;16MG/ML	A073199	001	Sep 11, 1992
TEVA PHARMS USA	80MG/ML;16MG/ML	A073303	001	Oct 31, 1991
WATSON LABS	80MG/ML;16MG/ML	A071556	001	Dec 29, 1987
WEST-WARD PHARMS INT	80MG/ML;16MG/ML	A070627	001	Dec 29, 1987
	80MG/ML;16MG/ML	A070628	001	Dec 29, 1987

SUSPENSION; ORAL

BACTRIM

+ SUN PHARM INDUSTRIES	200MG/5ML;40MG/5ML **	N017560	001	
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BACTRIM PEDIATRIC

SUN PHARM INDUSTRIES	200MG/5ML;40MG/5ML **	N017560	002	
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SEPTRA

MONARCH PHARMS	200MG/5ML;40MG/5ML **	N017598	001	
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SEPTRA GRAPE

MONARCH PHARMS	200MG/5ML;40MG/5ML **	N017598	002	Feb 12, 1986
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SULFAMETHOXAZOLE AND TRIMETHOPRIM

ANI PHARMS INC	200MG/5ML;40MG/5ML	A070028	001	Jun 02, 1987
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
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SULMEPRIM

USL PHARMA	200MG/5ML;40MG/5ML	A070063	001	Aug 01, 1986
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SULMEPRIM PEDIATRIC

USL PHARMA	200MG/5ML;40MG/5ML	A070064	001	Aug 01, 1986
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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
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COTRIM D.S.

TEVA	800MG;160MG	A070048	001	Mar 18, 1985
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SULFAMETHOPRIM

NOVEL LABS INC	400MG;80MG	A070022	001	Feb 15, 1985
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SULFAMETHOPRIM-DS

NOVEL LABS INC	800MG;160MG	A070032	001	Feb 15, 1985
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SULFAMETHOXAZOLE AND TRIMETHOPRIM

FOSUN PHARMA	400MG;80MG	A070889	001	Nov 13, 1986
	400MG;80MG	N018598	003	May 19, 1982
	800MG;160MG	A070890	001	Nov 13, 1986
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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

6-386(of 430)

** See List Footnote

SULFAMETHOXAZOLE; TRIMETHOPRIM

TABLET; ORAL

SULFAMETHOXAZOLE AND TRIMETHOPRIM

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
HERITAGE PHARMA	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	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

SULFANILAMIDE

ACP NIMBLE	15%	A088718 001	Sep 19, 1985
SUPPOSITORY; VAGINAL			
AVC			
MYLAN SPECIALITY LP	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	
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SULFASALAZINE

SUSPENSION; ORAL

AZULFIDINE

PHARMACIA AND UPJOHN	250MG/5ML	N018605 001	
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TABLET; ORAL

S.A.S.-500

SOLVAY	500MG	A083450 001	
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SULFASALAZINE

HERITAGE PHARMS INC	500MG	A080197 001	
SANDOZ	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

DISCONTINUED DRUG PRODUCT LIST

6-387(of 430)

** See List Footnote

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

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 INC 500MG

A080142 001

AUROLIFE PHARMA LLC 500MG

A085628 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

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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

6-388(of 430)

** See List Footnote

SULFOXONE SODIUM

TABLET, DELAYED RELEASE;ORAL
DIASONE SODIUM
ABBVIE

165MG

N006044 003

SULFUR

POWDER;TOPICAL
BENSULFOID

POYTHRESS

33.32%

N002918 001

SULINDAC

TABLET;ORAL
CLINORIL

+ MERCK

150MG **

N017911 001

+

200MG **

N017911 002

SULINDAC

ANI PHARMS INC

150MG

A072972 001 Feb 28, 1992

200MG

A072973 001 Feb 28, 1992

EPIC PHARMA LLC

150MG

A073262 002 Sep 06, 1991

200MG

A073262 001 Sep 06, 1991

FOSUN PHARMA

150MG

A072712 001 Aug 30, 1991

200MG

A072713 001 Aug 30, 1991

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

SUMATRIPTAN SUCCINATE

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

MYLAN LABS LTD

EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)

A203322 001 Apr 14, 2014

PAR STERILE PRODUCTS

EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)

A077871 001 Jul 09, 2009

SANDOZ INC

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

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 VENTURES LTD

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

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

DISCONTINUED DRUG PRODUCT LIST

6-389(of 430)

** See List Footnote

SUPROFEN

SOLUTION/DROPS;OPHTHALMIC

PROFENAL

ALCON

1%

N019387 001 Dec 23, 1988

SUTILAINS

OINTMENT;TOPICAL

TRAVASE

+ ABBOTT

82,000 UNITS/GM **

N012828 001

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

EQ 5MG BASE

A090402 001 Jul 01, 2010

TADALAFIL

TABLET;ORAL

TADALAFIL

WATSON LABS INC

2.5MG

A205885 001 Mar 29, 2019

5MG

A205885 002 Mar 29, 2019

10MG

A205885 003 Mar 29, 2019

20MG

A205885 004 Mar 29, 2019

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

MYLAN

0.4MG

A090408 001 Apr 27, 2010

TAPENTADOL HYDROCHLORIDE

SOLUTION;ORAL

NUCYNTA

+ COLLEGIUM PHARM INC

EQ 20MG BASE/ML

N203794 001 Oct 15, 2012

TAZAROTENE

CREAM;TOPICAL

TAZAROTENE

FOUGERA PHARMS INC

0.1%

A211175 001 Jan 28, 2019

DISCONTINUED DRUG PRODUCT LIST

6-390(of 430)

** See List Footnote

TECHNETIUM TC-99M ALBUMIN AGGREGATED

INJECTABLE; INJECTION			
TC 99M-LUNGAGGREGATE			
GE HEALTHCARE	5mCi/ML		N017848 001

TECHNETIUM TC-99M ALBUMIN AGGREGATED KIT

INJECTABLE; INJECTION			
A-N STANNOUS AGGREGATED ALBUMIN			
SYNCOR PHARMS	N/A		N017916 001
AN-MAA			
PHARMALUCENCE	N/A		N017792 001
LUNGAGGREGATE REAGENT			
GE HEALTHCARE	N/A		N017838 001
MACROTEC			
BRACCO	N/A		N017833 001
PULMOLITE			
+ JUBILANT DRAXIMAGE	N/A		N017776 001
TECHNESCAN MAA			
MALLINCKRODT	N/A		N017842 001
TECHNETIUM TC 99M MAA			
GE HEALTHCARE	N/A		N017773 001

TECHNETIUM TC-99M ALBUMIN COLLOID KIT

INJECTABLE; INJECTION			
MICROLITE			
PHARMALUCENCE	N/A		N018263 001 Mar 25, 1983

TECHNETIUM TC-99M ALBUMIN KIT

INJECTABLE; INJECTION			
TECHNETIUM TC 99M HSA			
GE HEALTHCARE	N/A		N017775 001

TECHNETIUM TC-99M ALBUMIN MICROSPHERES KIT

INJECTABLE; INJECTION			
INSTANT MICROSPHERES			
3M	N/A		N017832 001

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			
PHARMALUCENCE	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 FERMENTETATE KIT

INJECTABLE; INJECTION			
RENOTEC			
BRACCO	N/A		N017045 001

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

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 LIDOFEININ KIT

INJECTABLE; INJECTION

TECHNESCAN HIDA

DRAXIMAGE N/A

N018489 001 Oct 31, 1986

TECHNETIUM TC-99M MEDRONATE

INJECTABLE; INJECTION

DRAXIMAGE MDP-10

JUBILANT DRAXIMAGE N/A

N018035 001

TECHNETIUM TC-99M MEDRONATE KIT

INJECTABLE; INJECTION

AMERSCAN MDP KIT

GE HEALTHCARE N/A

N018335 001 Aug 05, 1982

MDP-BRACCO

CARDINAL HEALTH 414 N/A

N018107 001

OSTEOLITE

PHARMALUCENCE N/A

N017972 001

TECHNETIUM TC 99M MPI MDP

GE HEALTHCARE N/A

N018141 001

N/A

N018141 002 Jun 12, 1989

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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

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, 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

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

TECHNETIUM TC-99M TETROFOSMIN KIT

INJECTABLE;INJECTION

MYOVUE

+ GE HEALTHCARE N/A N020372 001 Feb 09, 1996

TEGASEROD MALEATE

TABLET;ORAL

ZELNORM

+ ALFASIGMA EQ 2MG BASE N021200 001 Jul 24, 2002

TELAPREVIR

TABLET;ORAL

INCIVEK

VERTEX PHARMS 375MG N201917 001 May 23, 2011

TELAVANCIN HYDROCHLORIDE

POWDER;INTRAVENOUS

VIBATIV

+ CUMBERLAND PHARMS 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

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

TELITHROMYCIN

TABLET; ORAL

KETEK

SANOFI AVENTIS US	300MG	N021144 002	Feb 09, 2005
	400MG	N021144 001	Apr 01, 2004

TELMISARTAN

TABLET; ORAL

TELMISARTAN

CIPLA	20MG	A078710 001	Jan 08, 2014
	40MG	A078710 002	Jan 08, 2014
	80MG	A078710 003	Jan 08, 2014
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
MICRO LABS	20MG	A207016 001	Oct 03, 2017
	40MG	A207016 002	Oct 03, 2017
	80MG	A207016 003	Oct 03, 2017

TEMAZEPAM

CAPSULE; ORAL

TEMAZ

QUANTUM PHARMICS	15MG	A070564 001	Oct 15, 1985
	30MG	A070547 001	Oct 15, 1985

TEMAZEPAM

DURAMED PHARMS BARR	15MG	A071708 001	Sep 29, 1988
	30MG	A071709 001	Sep 29, 1988
SUN PHARM INDUSTRIES	15MG	A071174 001	Jul 10, 1986
	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

TEMOZOLOMIDE

APOTEX INC	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
LANNETT CO INC	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
MYLAN	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

TENIPOSIDE

INJECTABLE; INJECTION

VUMON

+ HQ SPECLT PHARMA	10MG/ML	N020119 001	Jul 14, 1992
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DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

TENOFOVIR DISOPROXIL FUMARATE

TABLET;ORAL

TENOFOVIR DISOPROXIL FUMARATE

CHARTWELL	300MG	A206481	001	Jul 26, 2018
MYLAN	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

BEXIMCO PHARMS USA	EQ 1MG BASE	A075667	001	Jul 28, 2000
	EQ 2MG BASE	A075667	002	Jul 28, 2000
	EQ 5MG BASE	A075667	003	Jul 28, 2000
	EQ 10MG BASE	A075667	004	Jul 28, 2000
HERITAGE PHARMA	EQ 1MG BASE	A075614	002	Jan 30, 2001
	EQ 2MG BASE	A075614	001	Jan 30, 2001
	EQ 5MG BASE	A075614	003	Jan 30, 2001
	EQ 10MG BASE	A075614	004	Jan 30, 2001
MYLAN TECHNOLOGIES	EQ 1MG BASE	A075384	001	Dec 01, 2000
	EQ 2MG BASE	A075384	002	Dec 01, 2000
	EQ 5MG BASE	A075384	003	Dec 01, 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

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 1MG BASE	A074657	001	Apr 28, 2000
	EQ 2MG BASE	A074315	002	Dec 31, 1998
	EQ 2MG BASE	A074657	002	Apr 28, 2000
	EQ 5MG BASE	A074315	003	Dec 31, 1998
	EQ 5MG BASE	A074657	003	Apr 28, 2000
	EQ 10MG BASE	A074315	004	Dec 31, 1998
	EQ 10MG BASE	A074657	004	Apr 28, 2000
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

TERBINAFINE

GEL;TOPICAL

LAMISIL

GLAXOSMITHKLINE CONS	1%	N020846	001	Apr 29, 1998
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TERBINAFINE HYDROCHLORIDE

CREAM;TOPICAL

LAMISIL

NOVARTIS	1%	N020192	001	Dec 30, 1992
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GRANULE;ORAL

LAMISIL

+	NOVARTIS	EQ 125MG BASE/PACKET	N022071	001	Sep 28, 2007
+		EQ 187.5MG BASE/PACKET	N022071	002	Sep 28, 2007

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

TERBINAFINE HYDROCHLORIDE

SOLUTION; TOPICAL

LAMISIL

GLAXOSMITHKLINE CONS 1%

N020749 001 Oct 17, 1997

TABLET; ORAL

TERBINAFINE HYDROCHLORIDE

CHARTWELL EQ 250MG BASE

A078199 001 Jul 02, 2007

GEDEON RICHTER USA EQ 250MG BASE

A077065 001 Jul 02, 2007

MYLAN EQ 250MG BASE

A077136 001 Jul 02, 2007

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

DR REDDYS 1MG/ML

A076853 001 Jul 20, 2004

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

APOTEX 7MG

A209601 001 Nov 02, 2018

14MG

A209601 002 Nov 02, 2018

AUROBINDO PHARMA LTD 7MG

A209638 001 Oct 26, 2018

14MG

A209638 002 Oct 26, 2018

MSN 7MG

A209623 001 Apr 24, 2019

14MG

A209623 002 Apr 24, 2019

TEVA PHARMS USA 7MG

A209700 001 Sep 04, 2018

14MG

A209700 002 Sep 04, 2018

WATSON LABS TEVA 7MG

A209549 001 Jul 27, 2018

14MG

A209549 002 Jul 27, 2018

ZYDUS PHARMS 7MG

A209668 001 Nov 30, 2018

14MG

A209668 002 Nov 30, 2018

TERIPARATIDE

SOLUTION; SUBCUTANEOUS

FORTEO

LILLY 0.75MG/3ML (0.25MG/ML)

N021318 001 Nov 26, 2002

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

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
250MG

N016118 001

N016118 002

TESTOSTERONE

FILM, EXTENDED RELEASE; TRANSDERMAL

ANDRODERM

ALLERGAN 2.5MG/24HR
5MG/24HR

N020489 001 Sep 29, 1995

N020489 002 May 02, 1997

TESTODERM

ALZA 4MG/24HR
6MG/24HR

N019762 001 Oct 12, 1993

N019762 002 Oct 12, 1993

TESTODERM TTS

ALZA 5MG/24HR

N020791 001 Dec 18, 1997

GEL; TRANSDERMAL

TESTOSTERONE

ANI PHARMS INC 25MG/2.5GM PACKET
50MG/5GM PACKET
PAR PHARM 25MG/2.5GM PACKET
PERRIGO ISRAEL 25MG/2.5GM PACKET
50MG/5GM PACKET

N202763 001 Feb 14, 2012

N202763 002 Feb 14, 2012

A076744 001 May 23, 2007

N203098 002 Jan 31, 2013

N203098 003 Jan 31, 2013

GEL, METERED; TRANSDERMAL

TESTOSTERONE

PERRIGO ISRAEL 12.5MG/1.25GM ACTUATION

N203098 001 Jan 31, 2013

INJECTABLE; INJECTION

TESTOSTERONE

DR REDDYS 100MG/ML
WATSON LABS 25MG/ML
50MG/ML

A086417 001 Jul 07, 1983

A086420 001 May 10, 1983

A086419 001 Aug 23, 1983

SOLUTION, METERED; TRANSDERMAL

AXIRON

+ ELI LILLY AND CO 30MG/1.5ML ACTUATION **

N022504 001 Nov 23, 2010

TABLET, EXTENDED RELEASE; BUCCAL

STRIANT

+ AUXILIUM PHARMS LLC 30MG

N021543 001 Jun 19, 2003

TESTOSTERONE CYPIONATE

INJECTABLE; INJECTION

DEPO-TESTOSTERONE

PHARMACIA AND UPJOHN 50MG/ML

A085635 001

TESTOSTERONE CYPIONATE

MYLAN INSTITUTIONAL 200MG/ML
WATSON LABS 100MG/ML
100MG/ML
200MG/ML

A040652 001 Dec 11, 2006

A084401 001

A086029 001

A084401 002

TESTOSTERONE ENANTHATE

INJECTABLE; INJECTION

DELATESTRYL

ENDO PHARMS 200MG/ML
+

N009165 001

N009165 003

TESTOSTERONE ENANTHATE

MYLAN INSTITUTIONAL 200MG/ML
WATSON LABS 100MG/ML
100MG/ML
200MG/ML

A040647 001 Oct 05, 2009

A083667 001

A085599 001

A083667 002

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

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

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
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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

TETRACHEL

ANGUS	250MG	A060343	001
	500MG	A060343	003

TETRACYCLINE HYDROCHLORIDE

ABBOTT	250MG	A061802	001
	500MG	A061802	002
ANI PHARMS INC	250MG	A061471	001
ELKINS SINN	250MG	A060059	001
FERRANTE	125MG	A060173	001
	250MG	A060173	002
HEATHER	250MG	A061148	001
	500MG	A061148	002
HIKMA PHARMS	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
MYLAN	250MG	A060783	001
	500MG	A060783	002
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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

TETRACYCLINE HYDROCHLORIDE

CAPSULE; ORAL

TETRACYCLINE HYDROCHLORIDE

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

PFIPHARMECS	250MG	A060082 003	
	500MG	A060082 004	

FIBER, EXTENDED RELEASE; PERIODONTAL

ACTISITE

SCHIFF AND CO	12.7MG/FIBER	N050653 001	Mar 25, 1994
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FOR SOLUTION; TOPICAL

TOPICYCLINE

SHIRE	2.2MG/ML	N050493 001	
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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	
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SUSPENSION; ORAL

ACHROMYCIN V

LEDERLE	125MG/5ML	N050263 002	
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SUMYCIN

PAR PHARM	125MG/5ML	A060400 001	
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TETRACYCLINE HYDROCHLORIDE

ALPHARMA US PHARMS	125MG/5ML	A060633 001	
FERRANTE	125MG/5ML	A060174 001	
PROTER	125MG/5ML	A060446 001	
PUREPAC PHARM	125MG/5ML	A060291 001	

TETRACYN

PFIPHARMECS	125MG/5ML	A060095 001	
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TETRAMED

IVAX SUB TEVA PHARMS	125MG/5ML	A061468 001	
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SUSPENSION/DROPS; OPHTHALMIC

ACHROMYCIN

STORZ	1%	N050268 001	
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TABLET; ORAL

PANMYCIN

PHARMACIA AND UPJOHN	250MG	A061705 001	
	500MG	A061705 002	

SUMYCIN

PAR PHARM	50MG	A061147 003	
	100MG	A061147 002	
	250MG	A061147 001	
	500MG	A061147 004	

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	

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

TETRACYCLINE PHOSPHATE COMPLEX

CAPSULE;ORAL

TETREX

EQ 500MG HYDROCHLORIDE

N050212 003

THALLOUS CHLORIDE TL-201

INJECTABLE;INJECTION

THALLOUS CHLORIDE TL 201

BRACCO 1mCi/ML

N018548 001 Dec 30, 1982

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

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

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

THEOPHYLLINE

CAPSULE, EXTENDED RELEASE;ORAL

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	
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	
THEOPHYLLINE 0.08% AND DEXTROSE 5% IN PLASTIC CONTAINER					
B BRAUN	80MG/100ML	N019083	002	Nov 07, 1984	
THEOPHYLLINE 0.16% AND DEXTROSE 5% IN PLASTIC CONTAINER					
B BRAUN	160MG/100ML	N019083	003	Nov 07, 1984	
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 0.4% 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	
+ HOSPIRA INC	40MG/100ML	N019211	001	Dec 14, 1984	

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

6-401(of 430)

** See List Footnote

THEOPHYLLINE

INJECTABLE; INJECTION

THEOPHYLLINE IN DEXTROSE 5% IN PLASTIC CONTAINER

	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	
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SYRUP; ORAL

ACCURBRON

	SANOFI AVENTIS US	150MG/15ML	A088746 001	Nov 22, 1985
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AQUAPHYLLIN

	FERNDALE LABS	80MG/15ML	A087917 001	Jan 18, 1983
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SLO-PHYLLIN

	SANOFI AVENTIS US	80MG/15ML	A085187 001	
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THEOCLEAR-80

	CENT PHARMS	80MG/15ML	A087095 001	Mar 01, 1982
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THEOPHYLLINE

	ALPHARMA US PHARMS	80MG/15ML	A086001 001	
		150MG/15ML	A086545 001	

TABLET; ORAL

QUIBRON-T

	MONARCH PHARMS	300MG	A088656 001	Aug 22, 1985
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SLO-PHYLLIN

	SANOFI AVENTIS US	100MG	A085202 001	
		200MG	A085204 001	

THEOCLEAR-100

	CENT PHARMS	100MG	A085353 002	
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THEOCLEAR-200

	CENT PHARMS	200MG	A085353 001	
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THEOLAIR

	MEDICIS	125MG	A086399 001	
		250MG	A086399 002	

THEOPHYL-225

	ORTHO MCNEIL PHARM	225MG	A084726 001	
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TABLET, CHEWABLE; ORAL

THEOPHYL

	ORTHO MCNEIL PHARM	100MG	A086506 001	Sep 12, 1985
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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	
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QUIBRON-T/SR

	MONARCH PHARMS	300MG	A087563 001	Jun 21, 1983
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SUSTAIRE

	ROERIG	100MG	A085665 001	
		300MG	A085665 002	

T-PHYL

	PHARM RES ASSOC	200MG	A088253 001	Aug 17, 1983
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THEO-DUR

	SCHERING	100MG	A085328 001	
		200MG	A086998 001	
		300MG	A085328 002	
		450MG	A089131 001	Jun 25, 1986

THEOCHRON

	NOSTRUM PHARMS LLC	300MG	A087400 002	Jan 11, 1983
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Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

6-402(of 430)

** See List Footnote

THEOPHYLLINE

TABLET, EXTENDED RELEASE;ORAL

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

450MG

A081236 001 Nov 09, 1992

INWOOD LABS

450MG

A040034 001 Apr 28, 1995

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

THIABENDAZOLE

SUSPENSION;ORAL

MINTEZOL

MERCK SHARP DOHME

500MG/5ML

N016097 001

TABLET, CHEWABLE;ORAL

MINTEZOL

MERCK SHARP DOHME

500MG

N016096 001

THIAMINE HYDROCHLORIDE

INJECTABLE;INJECTION

BETALIN S

LILLY

100MG/ML

A080853 001

THIAMINE HYDROCHLORIDE

ABRAXIS PHARM

100MG/ML

A080509 001

AKORN

100MG/ML

A087968 001 Oct 01, 1982

BEL MAR

100MG/ML

A080718 001

200MG/ML

A080712 001

DELL LABS

100MG/ML

A083775 001

DR REDDYS

100MG/ML

A080571 001

200MG/ML

A080571 002

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

WEST-WARD PHARMS INT

100MG/ML

A080575 001

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

DISCONTINUED DRUG PRODUCT LIST

6-403(of 430)

** See List Footnote

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 INC

30MG/ML

A089602 001 Nov 09, 1987

HI TECH PHARMA

100MG/ML

A089603 001 Nov 09, 1987

HI TECH PHARMA

30MG/ML

A040125 001 Aug 16, 1996

PHARM ASSOC

100MG/ML

A040126 001 Aug 16, 1996

PHARM ASSOC

30MG/ML

A040187 001 Aug 28, 1997

SANDOZ

100MG/ML

A040213 001 May 29, 1998

SANDOZ

30MG/ML

A088307 001 Nov 23, 1983

WOCKHARDT

100MG/ML

A088308 001 Nov 23, 1983

WOCKHARDT

30MG/ML

A088258 001 Jul 25, 1983

WOCKHARDT

100MG/ML

A088227 001 Jul 05, 1983

THIORIDAZINE HYDROCHLORIDE INTENSOL

ROXANE

30MG/ML

A088941 001 Dec 16, 1985

ROXANE

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 INC

10MG

A088270 001 Apr 14, 1983

ANI PHARMS INC

10MG

A088493 001 May 17, 1985

ANI PHARMS INC

15MG

A088271 001 Apr 14, 1983

ANI PHARMS INC

25MG

A088272 001 Apr 14, 1983

ANI PHARMS INC

50MG

A088194 001 Apr 14, 1983

ANI PHARMS INC

100MG

A088273 001 Oct 03, 1983

ANI PHARMS INC

100MG

A088456 001 May 17, 1985

FOSUN PHARMA

10MG

A088131 001 Aug 30, 1983

FOSUN PHARMA

15MG

A088132 001 Aug 30, 1983

FOSUN PHARMA

25MG

A088133 001 Aug 30, 1983

FOSUN PHARMA

50MG

A088134 001 Aug 30, 1983

FOSUN PHARMA

100MG

A088135 001 Nov 20, 1984

FOSUN PHARMA

150MG

A088136 001 Sep 17, 1986

FOSUN PHARMA

200MG

A088137 001 Sep 17, 1986

HERITAGE PHARMA

10MG

A088476 001 Nov 08, 1983

HERITAGE PHARMA

25MG

A088478 001 Nov 08, 1983

HERITAGE PHARMA

50MG

A088479 001 Nov 08, 1983

HERITAGE PHARMA

100MG

A088736 001 Jul 24, 1984

MUTUAL PHARM

10MG

A088375 001 Nov 18, 1983

MUTUAL PHARM

25MG

A087264 001 Nov 18, 1983

MUTUAL PHARM

50MG

A088370 001 Nov 18, 1983

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

6-404(of 430)

** See List Footnote

THIORIDAZINE HYDROCHLORIDE

TABLET; ORAL

THIORIDAZINE HYDROCHLORIDE

	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
	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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

THIOTHIXENE

CAPSULE; ORAL

THIOTHIXENE

	2MG	A071885 001	Aug 12, 1987
	5MG	A071886 001	Aug 12, 1987
	10MG	A071887 001	Aug 12, 1987
	20MG	A072200 001	Dec 17, 1987
HERITAGE PHARMA	1MG	A070600 001	Jun 05, 1987
	2MG	A070601 001	Jun 05, 1987
	5MG	A070602 001	Jun 05, 1987
	10MG	A070603 001	Jun 05, 1987
SANDOZ	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	

THYROGLOBULIN

TABLET; ORAL

PROLOID

PARKE DAVIS	16MG	N002245 009	
	32MG	N002245 005	
	65MG	N002245 002	
	100MG	N002245 008	
	130MG	N002245 010	
	200MG	N002245 007	
	325MG	N002245 004	
THYROGLOBULIN			
IMPAX LABS	64.8MG	A080151 001	

THYROTROPIN

INJECTABLE; INJECTION

THYTROPAR

SANOFI AVENTIS US	10 IU/VIAL	N008682 001	
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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

TICAGRELOR

TABLET; ORAL

TICAGRELOR

AMNEAL PHARMS CO	90MG	A208531 001	Jan 23, 2019
WATSON LABS INC	60MG	A208390 001	Sep 04, 2018
	90MG	A208390 002	Sep 04, 2018

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

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
TICLOPIDINE HYDROCHLORIDE			
ACTAVIS ELIZABETH	250MG	A075253 001	Aug 20, 1999
MYLAN	250MG	A075161 001	Sep 13, 1999
	250MG	A075316 001	Nov 02, 1999
SUN PHARM INDS INC	250MG	A075526 001	Sep 26, 2002
WATSON LABS	250MG	A075309 001	Apr 26, 2000
YAOPHARMA CO LTD	250MG	A075318 001	Aug 20, 1999
	250MG	A075326 001	Aug 20, 1999

TILUDRONATE DISODIUM

TABLET; ORAL

SKELID

+ SANOFI AVENTIS US	EQ 200MG BASE **	N020707 001	Mar 07, 1997
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TIMOLOL MALEATE

SOLUTION/DROPS; OPHTHALMIC

TIMOLOL MALEATE

AKORN	EQ 0.25% BASE	A074465 001	Mar 25, 1997
	EQ 0.25% BASE	A074515 001	Mar 25, 1997
APOTEX INC	EQ 0.25% BASE	A075411 001	Sep 08, 2000
	EQ 0.5% BASE	A075412 001	Sep 08, 2000
FOUGERA	EQ 0.25% BASE	A074667 001	Mar 25, 1997
	EQ 0.5% BASE	A074668 001	Mar 25, 1997

TABLET; ORAL

BLOCADREN

+ MERCK	5MG **	N018017 001	
+	10MG **	N018017 002	
+	20MG **	N018017 004	

TIMOLOL MALEATE

ANI PHARMS INC	5MG	A072917 001	Jul 31, 1991
	10MG	A072918 001	Jul 31, 1991
	20MG	A072919 001	Jul 31, 1991
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
YAOPHARMA CO LTD	5MG	A072550 001	Apr 13, 1989
	10MG	A072551 001	Apr 13, 1989
	20MG	A072552 001	Apr 13, 1989

TINZAPARIN SODIUM

INJECTABLE; INJECTION

INNOHEP

LEO PHARMA AS	20,000 IU/ML	N020484 001	Jul 14, 2000
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DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

TIOCONAZOLE

CREAM; TOPICAL

TZ-3

PFIZER

1%

N018682 001 Feb 18, 1983

TIROFIBAN HYDROCHLORIDE

INJECTABLE; INJECTION

AGGRASTAT

MEDICURE

EQ 12.5MG BASE/50ML (EQ 0.25MG BASE/ML)

N020912 001 May 14, 1998

EQ 25MG BASE/500ML (EQ 0.05MG BASE/ML)

N020913 001 May 14, 1998

TIZANIDINE HYDROCHLORIDE

TABLET; ORAL

TIZANIDINE HYDROCHLORIDE

ANI PHARMS INC

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

MYLAN PHARMS INC

EQ 2MG BASE

A076282 001 Dec 16, 2003

EQ 4MG BASE

A076282 002 Dec 16, 2003

PAR PHARM INC

EQ 2MG BASE

A207170 001 Jan 26, 2017

EQ 4MG BASE

A207170 002 Jan 26, 2017

ZANAFLEX

+ COVIS PHARMA BV

EQ 2MG BASE **

N020397 002 Feb 04, 2000

TOBRAMYCIN

SOLUTION; INHALATION

TOBRAMYCIN

MYLAN

300MG/5ML

A209554 001 Oct 13, 2017

TEVA PHARMS USA

300MG/4ML

A210915 001 Jun 26, 2019

SOLUTION/DROPS; OPHTHALMIC

TOBRAMYCIN

ALCON PHARMS LTD

0.3%

A063176 001 May 25, 1994

APOTEX INC

0.3%

A065087 001 Feb 25, 2002

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

HOSPIRA

EQ 10MG BASE/ML

A063080 001 Apr 30, 1991

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

WATSON LABS INC

EQ 10MG BASE/ML

A062945 001 Aug 09, 1989

EQ 40MG BASE/ML

A062945 002 Aug 09, 1989

WEST-WARD PHARMS INT

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

TOBRAMYCIN SULFATE (PHARMACY BULK)

HOSPIRA

EQ 40MG BASE/ML **

A063116 001 May 18, 1992

DISCONTINUED DRUG PRODUCT LIST

6-408(of 430)

** See List Footnote

TOCAINIDE HYDROCHLORIDE

TABLET; ORAL

TONOCARD

ASTRAZENECA	400MG	N018257 001	Nov 09, 1984
	600MG	N018257 002	Nov 09, 1984

TOLAZAMIDE

TABLET; ORAL

TOLAZAMIDE

BARR	100MG	A070162 001	Jan 14, 1986
	250MG	A070163 001	Jan 14, 1986
	500MG	A070164 001	Jan 14, 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
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
	250MG	A070243 001	Aug 01, 1986
	250MG	A070514 001	Jan 09, 1986
	500MG	A070244 001	Aug 01, 1986
	500MG	A070515 001	Jan 09, 1986
YAOPHARMA CO LTD	100MG	A071633 001	Dec 09, 1987
	250MG	A070289 001	Mar 13, 1986
	500MG	A070290 001	Mar 13, 1986

TOLINASE

+ PHARMACIA AND UPJOHN	100MG **	N015500 002	
+	250MG **	N015500 004	
+	500MG **	N015500 005	

TOLAZOLINE HYDROCHLORIDE

INJECTABLE; INJECTION

PRISCOLINE

NOVARTIS	25MG/ML	N006403 005	Feb 22, 1985
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TOLBUTAMIDE

TABLET; ORAL

ORINASE

PHARMACIA AND UPJOHN	250MG **	N010670 002	
	500MG **	N010670 001	

TOLBUTAMIDE

ALRA	500MG	A086141 001	
ANI PHARMS INC	500MG	A087093 001	
ASCOT	500MG	A087541 001	Mar 01, 1983
BARR	500MG	A087121 001	
DAVA PHARMS INC	500MG	A086926 001	
PARKE DAVIS	500MG	A086047 001	
PUREPAC PHARM	500MG	A088950 001	Jun 17, 1985
SANDOZ	500MG	N012678 001	
SUPERPHARM	500MG	A088893 001	Nov 19, 1984
VANGARD	500MG	A087876 001	Apr 20, 1982
WATSON LABS	250MG	A089110 001	May 29, 1987
	500MG	A086109 001	
	500MG	A087318 001	
	500MG	A089111 001	May 29, 1987
YAOPHARMA CO LTD	500MG	A086574 001	

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

6-409(of 430)

** See List Footnote

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

TOLMETIN SODIUM

CAPSULE; ORAL

TOLECTIN DS

+ ORTHO MCNEIL JANSSEN EQ 400MG BASE

N018084 001

TOLMETIN SODIUM

ANI PHARMS INC EQ 400MG BASE

A073308 001 Jan 24, 1992

EQ 400MG BASE

A073392 001 Jan 24, 1992

EQ 400MG BASE

A073519 001 May 29, 1992

FOSUN PHARMA EQ 400MG BASE

A073462 001 Apr 30, 1992

SUN PHARM INDUSTRIES EQ 400MG BASE

A073311 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

TOLMETIN SODIUM

ACP NIMBLE EQ 600MG BASE

A074399 001 Mar 28, 1996

EQ 600MG BASE

A074729 001 Feb 27, 1997

ANI PHARMS INC EQ 600MG BASE

A073527 001 Jun 30, 1992

FOSUN PHARMA EQ 200MG BASE

A073588 001 Jul 31, 1992

EQ 600MG BASE

A074002 001 Sep 27, 1993

SUN PHARM INDUSTRIES EQ 200MG BASE

A073310 001 Nov 27, 1991

TOLTERODINE TARTRATE

TABLET; ORAL

TOLTERODINE TARTRATE

APOTEX CORP 1MG

A200164 001 Sep 25, 2012

2MG

A200164 002 Sep 25, 2012

MYLAN PHARMS INC 1MG

A202641 001 Nov 27, 2012

2MG

A202641 002 Nov 27, 2012

TOLVAPTAN

TABLET; ORAL

SAMSCA

+ OTSUKA AMERICA PHARM 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

FOSUN PHARMA 15MG

A079206 001 Oct 14, 2009

25MG

A079206 002 Oct 14, 2009

MYLAN 15MG

A078418 001 Oct 14, 2009

25MG

A078418 002 Oct 14, 2009

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

BARR 25MG

A076315 001 Mar 27, 2009

100MG

A076315 002 Mar 27, 2009

200MG

A076315 003 Mar 27, 2009

HIKMA PHARMS 25MG

A091185 001 Nov 25, 2013

50MG

A091185 002 Nov 25, 2013

100MG

A091185 003 Nov 25, 2013

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

TOPIRAMATETABLET; ORAL
TOPIRAMATE

	200MG	A091185 004	Nov 25, 2013
LUPIN	25MG	A078410 001	Sep 11, 2013
	50MG	A078410 002	Sep 11, 2013
	100MG	A078410 003	Sep 11, 2013
	200MG	A078410 004	Sep 11, 2013
MYLAN	25MG	A076314 001	Mar 27, 2009
	50MG	A076314 002	Mar 27, 2009
	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
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 USA	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
MYLAN LABS LTD	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

TORSEMIDE

INJECTABLE; INJECTION

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
WEST-WARD PHARMS INT	20MG/2ML (10MG/ML)	A078007 001	Jun 11, 2008
	50MG/5ML (10MG/ML)	A078007 002	Jun 11, 2008

TABLET; ORAL

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

TRAMADOL HYDROCHLORIDE

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
FOSUN PHARMA	50MG	A075968 001	Jun 25, 2002
IVAX SUB TEVA PHARMS	50MG	A075963 001	Jul 03, 2002
MYLAN PHARMS INC	50MG	A075980 001	Nov 21, 2002
NORTHSTAR HLTHCARE	50MG	A078935 001	May 26, 2010
POLYGEN PHARMS	50MG	A206706 001	Jul 02, 2019
SPECGX LLC	50MG	A075983 001	Jun 25, 2002
SUN PHARM INDUSTRIES	50MG	A076100 001	Jun 20, 2002

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

TRAMADOL HYDROCHLORIDE

TABLET; ORAL

TRAMADOL HYDROCHLORIDE					
WATSON LABS	50MG		A075962	001	Jun 24, 2002
ULTRAM					
JANSSEN PHARMS	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					
AUROBINDO PHARMA LTD	100MG		A204421	001	Oct 20, 2015
	200MG		A204421	002	Oct 20, 2015
	300MG		A204421	003	Oct 20, 2015
PAR PHARM INC	100MG		A078783	001	Nov 13, 2009
	200MG		A078783	002	Nov 13, 2009
	300MG		A078783	003	Sep 20, 2011
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

TRANDOLAPRIL

TABLET; ORAL

MAVIK					
+ ABBVIE	1MG **		N020528	001	Apr 26, 1996
+	2MG **		N020528	002	Apr 26, 1996
+	4MG **		N020528	003	Apr 26, 1996
TRANDOLAPRIL					
CIPLA	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
MYLAN	1MG		A078346	001	Apr 28, 2008
	2MG		A078346	002	Apr 28, 2008
	4MG		A078346	003	Apr 28, 2008
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

TRANEXAMIC ACID

INJECTABLE; INJECTION

TRANEXAMIC ACID					
CAPLIN	100MG/ML		A212360	001	Jul 17, 2019
VIRTUS PHARMS	100MG/ML		A202755	001	Feb 25, 2016
TABLET; ORAL					
CYKLOKAPRON					
PHARMACIA AND UPJOHN	500MG		N019280	001	Dec 30, 1986

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

TRANEXAMIC ACID

TABLET;ORAL

TRANEXAMIC ACID

ANI PHARMS INC 650MG A203256 001 Jul 25, 2016

TRAVOPROST

SOLUTION/DROPS;OPHTHALMIC

IZBA

+ NOVARTIS 0.003% ** N204822 001 May 15, 2014

TRAVATAN

+ ALCON PHARMS LTD 0.004% ** N021257 001 Mar 16, 2001

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

AM THERAP

50MG A071139 001 Oct 29, 1986

100MG A071140 001 Oct 29, 1986

AUROLIFE PHARMA LLC

50MG A072484 001 Apr 30, 1990

FOSUN PHARMA

100MG A072483 001 Apr 30, 1990

MYLAN

50MG A071405 001 Feb 27, 1991

100MG A071406 001 Feb 27, 1991

MYLAN PHARMS INC

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

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

TRETINOIN

CAPSULE;ORAL

VESANOID

+ CHEPLAPHARM 10MG ** N020438 001 Nov 22, 1995

CREAM;TOPICAL

TRETINOIN

ALLERGAN 0.0375% A090098 001 Mar 22, 2010

0.075% A202209 001 Oct 11, 2012

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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

TRIAMCINOLONE

TABLET; ORAL

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

IMPAX LABS	4MG	A084340 001
IVAX SUB TEVA PHARMS	4MG	A083750 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
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AEROSOL, METERED; NASAL

NASACORT

SANOVI AVENTIS US	0.055MG/INH	N019798 001	Jul 11, 1991
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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
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KENALOG-H

DELCOR ASSET CORP	0.1%	A086240 001
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TRACET

TEVA	0.025%	A084908 001
	0.1%	A084908 002
	0.5%	A084908 003

TRACORT

SOLVAY	0.1%	A087113 001
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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
PHARMADERM	0.025%	A087990 001	Jul 07, 1983
	0.1%	A087991 001	Jul 07, 1983
	0.5%	A087992 001	Jul 07, 1983

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

TRIAMCINOLONE ACETONIDE

CREAM;TOPICAL

TRIAMCINOLONE ACETONIDE

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	
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INJECTABLE; INJECTION

TRIAMCINOLONE ACETONIDE

PARNELL	3MG/ML	N019503 001	Oct 16, 1987
SANDOZ INC	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
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LOTION;TOPICAL

KENALOG

DELacor 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

OINTMENT;TOPICAL

ARISTOCORT

ASTELLAS	0.1%	A080750 004	
	0.5% **	A080745 002	

ARISTOCORT A

ASTELLAS	0.1%	A080750 003	
	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

DELacor ASSET CORP	0.5% **	A083944 001	
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TRIAMCINOLONE ACETONIDE

ACTAVIS MID ATLANTIC	0.1%	A087799 001	Jun 07, 1982
ALPHARMA US PHARMS	0.5%	A089913 001	Dec 23, 1988
MORTON GROVE	0.025%	A088090 001	Sep 01, 1983
	0.1%	A088091 001	Sep 01, 1983
	0.5%	A088092 001	Sep 01, 1983
+ MYLAN	0.025% **	N011600 003	
+	0.1% **	N011600 001	
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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

6-415(of 430)

** See List Footnote

TRIAMCINOLONE ACETONIDE

OINTMENT; TOPICAL

TRIAMCINOLONE ACETONIDE

0.1%	A087902 001	Dec 27, 1982
0.5%	A040386 001	Jun 05, 2001

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

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 ISRAEL

0.055MG/SPRAY

A078104 001 Jul 30, 2009

TRIAMCINOLONE DIACETATE

INJECTABLE; INJECTION

ARISTOCORT

FOSUN PHARMA

25MG/ML

N011685 003

+

TRIAMCINOLONE DIACETATE

AKORN

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

TRIAZOLAM

TABLET; ORAL

HALCION

PHARMACIA AND UPJOHN

0.5MG

N017892 002 Nov 15, 1982

TRIAZOLAM

MYLAN PHARMS INC

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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

6-416(of 430)

** See List Footnote

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

TRIFLUOPERAZINE HYDROCHLORIDE

CONCENTRATE; ORAL

STELAZINE

+ GLAXOSMITHKLINE EQ 10MG BASE/ML ** N011552 006

TRIFLUOPERAZINE HYDROCHLORIDE

FOSUN PHARMA 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

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

INVATECH 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

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

DISCONTINUED DRUG PRODUCT LIST

6-417(of 430)

** See List Footnote

TRIHEXYPHENIDYL HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE;ORAL

ARTANE

LEDERLE	5MG	N006773	010
	5MG	N012947	001

ELIXIR;ORAL

ARTANE

LEDERLE	2MG/5ML	N006773	009
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TRIHEXYPHENIDYL HYDROCHLORIDE

PHARM VENTURES	2MG/5ML	A089514	001	Apr 07, 1989
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TABLET;ORAL

ARTANE

+ LEDERLE	2MG **	N006773	005
+ LEDERLE	5MG **	N006773	003

TREMIM

SCHERING	2MG	A080381	001
	5MG	A080381	003

TRIHEXYPHENIDYL HYDROCHLORIDE

HIKMA PHARMS	2MG	A040337	002	Feb 16, 2000
	5MG	A040337	001	Feb 16, 2000

NYLOS	5MG	A085622	001
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VANGARD	2MG	A088035	001	Jul 30, 1982
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WATSON LABS	2MG	A040184	001	Feb 06, 1998
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	2MG	A085117	001
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	5MG	A040184	002	Feb 06, 1998
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	5MG	A085105	001
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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
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SYRUP;ORAL

TEMARIL

ALLERGAN HERBERT	EQ 2.5MG BASE/5ML	N011316	003
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TRIMEPRAZINE TARTRATE

ALPHARMA US PHARMS	EQ 2.5MG BASE/5ML	A085015	001	Feb 18, 1982
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MORTON GROVE	EQ 2.5MG BASE/5ML	A088285	001	Apr 11, 1985
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TABLET;ORAL

TEMARIL

ALLERGAN HERBERT	EQ 2.5MG BASE	N011316	001
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TRIMETHADIONE

CAPSULE;ORAL

TRIDIONE

ABBVIE	300MG	N005856	005
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SOLUTION;ORAL

TRIDIONE

ABBVIE	200MG/5ML	N005856	002
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TABLET;ORAL

TRIDIONE

+ ABBVIE	150MG	N005856	009
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TRIMETHAPHAN CAMSYLATE

INJECTABLE; INJECTION

ARFONAD

ROCHE	50MG/ML	N008983	001
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TRIMETHOBENZAMIDE HYDROCHLORIDE

INJECTABLE; INJECTION

TRIMETHOBENZAMIDE HYDROCHLORIDE

AM REGENT	100MG/ML	A091330	001	Mar 08, 2011
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HOSPIRA	100MG/ML	A088804	001	Apr 03, 1987
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SMITH AND NEPHEW	100MG/ML	A088960	001	Apr 04, 1986
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	100MG/ML	A089043	001	Apr 04, 1986
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SOLOPAK	100MG/ML	A089094	001	Apr 04, 1986
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WATSON LABS	100MG/ML	A086577	001	Oct 19, 1982
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Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

6-418(of 430)

** See List Footnote

TRIMETHOBENZAMIDE HYDROCHLORIDE

INJECTABLE; INJECTION

TRIMETHOBENZAMIDE HYDROCHLORIDE

100MG/ML

A087939 001 Dec 28, 1982

TRIMETHOBENZAMIDE HYDROCHLORIDE PRESERVATIVE FREE

AM REGENT

100MG/ML

A091329 001 Mar 08, 2011

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

ALLEGIS

EQ 25MG BASE/5ML

N074374 001 Jun 23, 1995

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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

6-419(of 430)

** See List Footnote

TRIPLENNAMINE HYDROCHLORIDETABLET, EXTENDED RELEASE;ORAL
PBZ-SR

NOVARTIS	50MG	N010533	002
	100MG	N010533	001

TRIPLE SULFA (SULFABENZAMIDE;SULFACETAMIDE;SULFATHIAZOLE)

CREAM;VAGINAL

GYNE-SULF

ACP NIMBLE	3.7%;2.86%;3.42%	A088607	001	Jun 09, 1986
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SULTRIN

ORTHO MCNEIL PHARM	3.7%;2.86%;3.42%	N005794	001
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TRIPLE SULFA

ALPHARMA US PHARMS	3.7%;2.86%;3.42%	A087864	001	Sep 01, 1982
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FOUGERA	3.7%;2.86%;3.42%	A086424	001
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PERRIGO NEW YORK	3.7%;2.86%;3.42%	A087285	001	Nov 15, 1982
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TRYSUL

SAVAGE LABS	3.7%;2.86%;3.42%	A087887	001	Jul 23, 1982
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VAGILIA

ACP NIMBLE	3.7%;2.86%;3.42%	A088821	001	Nov 09, 1987
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TABLET;VAGINAL

SULTRIN

ORTHO MCNEIL PHARM	184MG;143.75MG;172.5MG	N005794	002
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TRIPLE SULFA

FOUGERA	184MG;143.75MG;172.5MG	A088463	001	Jan 03, 1985
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PHARMADERM	184MG;143.75MG;172.5MG	A088462	001	Jan 03, 1985
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TRIPROLIDINE HYDROCHLORIDE

SYRUP;ORAL

ACTIDIL

GLAXOSMITHKLINE	1.25MG/5ML	N011496	002	Jul 01, 1983
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MYIDYL

USL PHARMA	1.25MG/5ML	A087963	001	Jan 18, 1983
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TRIPROLIDINE HYDROCHLORIDE

ALPHARMA US PHARMS	1.25MG/5ML	A085940	001
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HALSEY	1.25MG/5ML	A088735	001	Jan 17, 1985
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PHARM ASSOC	1.25MG/5ML	A087514	001	Feb 10, 1982
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TABLET;ORAL

ACTIDIL

GLAXOSMITHKLINE	2.5MG	N011110	002	Jul 01, 1983
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TRIPROLIDINE HYDROCHLORIDE

VITARINE	2.5MG	A085610	001
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WATSON LABS	2.5MG	A085094	001
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TRISULFAPYRIMIDINES (SULFADIAZINE;SULFAMERAZINE;SULFAMETHAZINE)

SUSPENSION;ORAL

LANTRISUL

LANNETT	167MG/5ML;167MG/5ML;167MG/5ML	A080123	002
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NEOTRIZINE

LILLY	167MG/5ML;167MG/5ML;167MG/5ML	N006317	012
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SULFALOID

FOREST PHARMS	167MG/5ML;167MG/5ML;167MG/5ML	A080100	001
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SULFOSE

WYETH AYERST	167MG/5ML;167MG/5ML;167MG/5ML	A080013	002
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TERFONYL

BRISTOL MYERS SQUIBB	167MG/5ML;167MG/5ML;167MG/5ML	N006904	002
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TRIPLE SULFA

ALPHARMA US PHARMS	167MG/5ML;167MG/5ML;167MG/5ML	A080280	001
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TRIPLE SULFAS

LEDERLE	167MG/5ML;167MG/5ML;167MG/5ML	N006920	003
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TABLET;ORAL

NEOTRIZINE

LILLY	167MG;167MG;167MG	N006317	011
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SULFA-TRIPLE #2

IMPAX LABS	167MG;167MG;167MG	A080079	001
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SULFALOID

FOREST PHARMS	167MG;167MG;167MG	A080099	001
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SULFOSE

WYETH AYERST	167MG;167MG;167MG	A080013	001
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Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

6-420(of 430)

** See List Footnote

TRISULFAPYRIMIDINES (SULFADIAZINE;SULFAMERAZINE;SULFAMETHAZINE)

TABLET;ORAL

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

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

TROMETHAMINE

INJECTABLE;INJECTION

THAM

+ HOSPIRA 3.6GM/100ML ** N013025 002

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

AKORN 1% A088447 001 Aug 28, 1985

ALCON PHARMS LTD 1% A089172 001 Dec 28, 1990

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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

6-421(of 430)

** See List Footnote

TUBOCURARINE CHLORIDE

INJECTABLE; INJECTION

TUBOCURARINE CHLORIDE

BRISTOL MYERS SQUIBB	3MG/ML	N005657	001
HOSPIRA	3MG/ML	N006095	001
LILLY	3MG/ML	N006325	001

TYROPANOATE SODIUM

CAPSULE; ORAL

BILOPAQUE

GE HEALTHCARE	750MG	N013731	001
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UNOPROSTONE ISOPROPYL

SOLUTION/DROPS; OPHTHALMIC

RESCULA

+ SUCAMPO PHARMA LLC	0.15% **	N021214	001	Aug 03, 2000
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URACIL MUSTARD

CAPSULE; ORAL

URACIL MUSTARD

SHIRE	1MG	N012892	001
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UREA

INJECTABLE; INJECTION

STERILE UREA

HOSPIRA	40GM/VIAL	N017698	001
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UREAPHIL

HOSPIRA	40GM/VIAL	N012154	001
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UREA C-13

FOR SOLUTION; ORAL

BREATHTEK UBT FOR H-PYLORI

OTSUKA AMERICA	EQ 75MG/POUCH	N020586	002	May 10, 2001
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HELICOSOL

METABOLIC SOLUTIONS	125MG/VIAL	N021092	001	Dec 17, 1999
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MERETEK UBT KIT (W/ PRANACTIN)

OTSUKA AMERICA	125MG/VIAL	N020586	001	Sep 17, 1996
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PYLORI-CHEK BREATH TEST

DXS DEVICES	100MG/VIAL	N020900	001	Feb 04, 1999
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UROFOLLITROPIN

INJECTABLE; INTRAMUSCULAR

METRODIN

SERONO	75 IU/AMP	N019415	002	Sep 18, 1986
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	150 IU/AMP	N019415	003	Sep 18, 1986
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INJECTABLE; INTRAMUSCULAR, SUBCUTANEOUS

BRAVELLE

+ FERRING	75 IU/VIAL	N021289	001	May 06, 2002
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INJECTABLE; SUBCUTANEOUS

FERTINEX

SERONO	75 IU/AMP	N019415	005	Aug 23, 1996
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	150 IU/AMP	N019415	004	Aug 23, 1996
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UROKINASE

INJECTABLE; INJECTION

KINLYTIC

MICROBIX BIOSYSTEMS	5,000 IU/VIAL	N021846	003
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	9,000 IU/VIAL	N021846	002
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	250,000 IU/VIAL	N021846	001
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URSODIOL

CAPSULE; ORAL

ACTIGALL

ALLERGAN	150MG	N019594	001	Dec 31, 1987
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URSODIOL

IMPAX LABS INC	300MG	A077895	001	Jul 27, 2006
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TABLET; ORAL

URSODIOL

TEVA PHARMS USA	250MG	A079184	001	May 13, 2009
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	500MG	A079184	002	May 13, 2009
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DISCONTINUED DRUG PRODUCT LIST

6-422(of 430)

** See List Footnote

VALACYCLOVIR HYDROCHLORIDE

TABLET; ORAL

VALACYCLOVIR HYDROCHLORIDE

MYLAN	EQ 500MG BASE	A078070 001	May 24, 2010
	EQ 1GM BASE	A078070 002	May 24, 2010
TEVA PHARMS	EQ 500MG BASE	A077655 001	May 24, 2010
	EQ 1GM BASE	A077655 002	May 24, 2010
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

VALPROATE SODIUM

INJECTABLE; INJECTION

DEPACON

+ ABBVIE	EQ 100MG BASE/ML	N020593 001	Dec 30, 1996
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VALPROIC ACID

CAPSULE; ORAL

DEPAKENE

+ ABBVIE	250MG	N018081 001	
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VALPROIC ACID

PAR PHARM	250MG	A070431 001	Feb 28, 1986
SCHERER RP	250MG	A070195 001	Jul 02, 1987
UPSHER SMITH LABS	250MG	A070631 001	Jun 11, 1987

CAPSULE, DELAYED RELEASE; ORAL

STAVZOR

+ BIONPHARMA INC	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	
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VALPROIC ACID

ANI PHARMS INC	250MG/5ML	A073178 001	Aug 25, 1992
APOTEX	250MG/5ML	A077105 001	Jul 29, 2005

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

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
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 HYDROCHLORIDE

CAPSULE; ORAL

VANCOMYCIN HYDROCHLORIDE

FRESENIUS KABI USA	EQ 125MG BASE	A065453 001	Jun 18, 2012
	EQ 250MG BASE	A065453 002	Jun 18, 2012

FOR SOLUTION; ORAL

VANCOCIN HYDROCHLORIDE

ANI PHARMS INC	EQ 500MG BASE/6ML	A061667 001	
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VANCOLED

LEDERLE	EQ 250MG BASE/5ML	A063321 002	Oct 15, 1993
	EQ 500MG BASE/6ML	A063321 003	Oct 15, 1993

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

VANCOMYCIN HYDROCHLORIDE

INJECTABLE; INJECTION

VANCOCIN HYDROCHLORIDE

ANI PHARMS INC

EQ 500MG BASE/VIAL **
 EQ 500MG BASE/VIAL
 EQ 500MG BASE/VIAL
 EQ 500MG BASE/VIAL **
 EQ 1GM BASE/VIAL **
 EQ 1GM BASE/VIAL
 EQ 1GM BASE/VIAL
 EQ 1GM BASE/VIAL **
 EQ 10GM BASE/VIAL **

A060180 001
 A062476 001 Mar 15, 1984
 A062716 001 Mar 13, 1987
 A062812 001 Nov 17, 1987
 A060180 002 Mar 21, 1986
 A062476 002 Mar 21, 1986
 A062716 002 Mar 13, 1987
 A062812 002 Nov 17, 1987
 A062812 003 Nov 17, 1987

VANCOLED

WEST-WARD PHARMS INT

EQ 500MG BASE/VIAL **
 EQ 1GM BASE/VIAL **
 EQ 2GM BASE/VIAL **
 EQ 5GM BASE/VIAL **
 EQ 10GM BASE/VIAL **

A062682 001 Jul 22, 1986
 A062682 002 Mar 30, 1988
 A062682 003 May 11, 1988
 A062682 004 May 11, 1988
 A062682 005 May 11, 1988

VANCOMYCIN HYDROCHLORIDE

EMCURE PHARMS LTD

EQ 500MG BASE/VIAL
 EQ 1GM BASE/VIAL
 EQ 10GM BASE/VIAL
 EQ 5GM BASE/VIAL

A202275 001 Oct 31, 2013
 A202275 002 Oct 31, 2013
 A202464 001 Oct 09, 2013
 A202274 001 Oct 31, 2013

HAINAN POLY PHARM

EQ 500MG BASE/VIAL
 EQ 1GM BASE/VIAL

A212332 001 Jun 12, 2019
 A212332 002 Jun 12, 2019

MYLAN LABS LTD

EQ 10GM BASE/VIAL

A091469 001 Jul 01, 2011

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

WEST-WARD PHARMS INT

EQ 500MG BASE/VIAL

A062879 001 Aug 02, 1988

EQ 1GM BASE/VIAL

A062879 002 Aug 02, 1988

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

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

VARDENAFIL HYDROCHLORIDE

TABLET; ORAL

LEVITRA

+ BAYER HLTHCARE

2.5MG **

N021400 003 Aug 19, 2003

VARDENAFIL HYDROCHLORIDE

AMNEAL PHARMS CO

5MG

A210738 001 Oct 31, 2018

10MG

A210738 002 Oct 31, 2018

20MG

A210738 003 Oct 31, 2018

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

HOSPIRA

4MG/VIAL

A075558 001 Sep 11, 2001

WATSON LABS

10MG/VIAL

A074334 001 Aug 31, 1995

20MG/VIAL

A074334 002 Aug 31, 1995

WEST-WARD PHARMS INT

10MG/VIAL

A075218 001 Aug 23, 1999

20MG/VIAL

A075218 002 Aug 23, 1999

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

VELAGLUCERASE ALFAPOWDER; INTRAVENOUS
VPRIV

SHIRE HUMAN GENETIC 200 UNITS/VIAL N022575 002 Feb 26, 2010

VENLAFAXINE HYDROCHLORIDECAPSULE, EXTENDED RELEASE; ORAL
EFFEXOR XR

WYETH PHARMS	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
MYLAN	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

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

FOSUN PHARMA	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
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

VERAPAMIL HYDROCHLORIDECAPSULE, EXTENDED RELEASE; ORAL
VERAPAMIL HYDROCHLORIDE

MYLAN	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
	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

TABLET; ORAL

CALAN

GD SEARLE LLC	40MG	N018817 003	Feb 23, 1988
+	80MG	N018817 001	Sep 10, 1984

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

VERAPAMIL HYDROCHLORIDE

TABLET; ORAL

CALAN

+	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

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
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SUN PHARM INDUSTRIES

80MG	A071489 002	Jan 13, 1988
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120MG	A071489 001	Jan 13, 1988
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WARNER CHILCOTT

80MG	A070340 001	Sep 24, 1986
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120MG	A070341 001	Sep 24, 1986
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WATSON LABS

40MG	A072799 001	Apr 28, 1989
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40MG	A072923 001	Jun 29, 1993
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80MG	A070855 001	Sep 24, 1986
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80MG	A071366 001	Oct 01, 1986
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120MG	A070856 001	Sep 24, 1986
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120MG	A071367 001	Oct 01, 1986
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YAOPHARMA CO LTD

40MG	A073168 001	Jul 31, 1992
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80MG	A071423 001	May 24, 1988
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120MG	A071424 001	May 25, 1988
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TABLET, EXTENDED RELEASE; ORAL

CALAN SR

+	PFIZER	180MG **	N019152 002	Dec 15, 1989
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COVERA-HS

GD SEARLE LLC

180MG	N020552 001	Feb 26, 1996
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240MG	N020552 002	Feb 26, 1996
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VERAPAMIL HYDROCHLORIDE

APOTEX CORP

120MG	A200878 001	Apr 20, 2012
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180MG	A200878 002	Apr 20, 2012
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240MG	A200878 003	Apr 20, 2012
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PLIVA

240MG	A072922 001	Mar 01, 1996
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VERATRUM VIRIDE ROOT

TABLET; ORAL

VERTAVIS

MEDPOINTE PHARM HLC

130CSR UNIT	N005691 002	
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VIDARABINE

INJECTABLE; INJECTION

VIRA-A

PARKEDALE

EQ 187.4MG BASE/ML	N050523 001	
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OINTMENT; OPHTHALMIC

VIRA-A

PARKEDALE

3%	N050486 001	
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VINBLASTINE SULFATE

INJECTABLE; INJECTION

VELBAN

+ LILLY

10MG/VIAL	N012665 001	
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VINBLASTINE SULFATE

ABRAXIS PHARM

10MG/VIAL	A089011 001	Nov 18, 1985
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HOSPIRA

10MG/VIAL	A089565 001	Aug 18, 1987
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VINCRISTINE SULFATE

INJECTABLE; INJECTION

ONCOVIN

LILLY

1MG/VIAL	N014103 001	
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1MG/ML	N014103 003	Mar 07, 1984
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5MG/VIAL	N014103 002	
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VINCASAR PFS

TEVA PARENTERAL

1MG/ML	A071426 001	Jul 17, 1987
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Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

VINCRIStINE SULFATE

INJECTABLE; INJECTION

VINCRESX

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

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

HOSPIRA

EQ 10MG BASE/ML

A076827 001 Jun 02, 2005

MYLAN LABS LTD

EQ 10MG BASE/ML

A200148 001 Aug 31, 2012

NOVAST LABS

EQ 10MG BASE/ML

A208997 001 Aug 05, 2019

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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

VITAMIN A PALMITATE

CAPSULE;ORAL

VITAMIN A PALMITATE

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

VORICONAZOLE

FOR SUSPENSION;ORAL

VORICONAZOLE

MYLAN PHARMS INC

200MG/5ML

A202361 001 May 28, 2013

TABLET;ORAL

VORICONAZOLE

TEVA PHARMS

50MG

A091658 001 Apr 06, 2012

200MG

A091658 002 Apr 06, 2012

VORTIOXETINE HYDROBROMIDE

TABLET;ORAL

TRINTELLIX

+ TAKEDA PHARMS USA

EQ 15MG BASE **

N204447 003 Sep 30, 2013

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

PANWARFIN

ABBOTT

2MG

N017020 001

2.5MG

N017020 002

5MG

N017020 003

7.5MG

N017020 004

10MG

N017020 005

WARFARIN SODIUM

MYLAN

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

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

YAOPHARMA CO LTD

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

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

WARFARIN SODIUM

TABLET; ORAL

WARFARIN SODIUM

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

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
5-100 CI/CYLINDER	N017550 001
0.25-5 CI/AMP	N017550 003

GEN ELECTRIC

XENON XE 133-V.S.S.

GE HEALTHCARE

10mCi/VIAL	N017687 001
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INJECTABLE; INJECTION

XENON XE 133

GE HEALTHCARE

1.3-1.7 CI/AMP	N017256 001
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LANTHEUS MEDCL

6.3mCi/ML	N017283 001
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SOLUTION; INHALATION, INJECTION

XENEISOL

MALLINCKRODT

18-25mCi/AMP	N017262 002
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XYLOSE

POWDER; ORAL

XYLO-PFAN

SAVAGE LABS

25GM/BOT	N017605 001
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XYLOSE

LYNE

25GM/BOT	N018856 001	Mar 26, 1987
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ZALCITABINE

TABLET; ORAL

HIVID

ROCHE

0.375MG	N020199 001	Jun 19, 1992
0.75MG	N020199 002	Jun 19, 1992

ZALEPLON

CAPSULE; ORAL

ZALEPLON

MYLAN

5MG	A077238 001	Jun 06, 2008
10MG	A077238 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 THERAPS LLC

200MCG/2ML (100MCG/ML)	N021060 003	Dec 28, 2004
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ZIDOVUDINE

INJECTABLE; INJECTION

ZIDOVUDINE

AM REGENT

10MG/ML	A091457 001	May 06, 2010
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LIAONING CHENGDA

10MG/ML	A204538 001	Nov 26, 2013
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TABLET; ORAL

RETROVIR

VIIV HLTHCARE

200MG	N020518 001	Dec 19, 1995
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+

300MG **	N020518 002	Oct 04, 1996
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ZIDOVUDINE

AUROBINDO PHARMA

60MG	N022294 001	Jul 23, 2009
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HEC PHARM

300MG	A202058 001	Oct 07, 2011
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Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

ZIDOVUDINE

TABLET;ORAL

ZIDOVUDINE

MATRIX LABS LTD	100MG	N200732 001	Feb 23, 2011
RANBAXY LABS LTD	300MG	A077327 001	Sep 19, 2005

ZILEUTON

TABLET;ORAL

ZYFLO

CHIESI USA INC	300MG	N020471 001	Dec 09, 1996
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ZINC SULFATE

INJECTABLE; INJECTION

ZINC SULFATE

ABRAXIS PHARM	EQ 1MG ZINC/ML	N019229 002	May 05, 1987
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ZIPRASIDONE HYDROCHLORIDE

CAPSULE;ORAL

ZIPRASIDONE HYDROCHLORIDE

CADILA	EQ 20MG BASE	A208988 001	Aug 22, 2017
	EQ 40MG BASE	A208988 002	Aug 22, 2017
	EQ 60MG BASE	A208988 003	Aug 22, 2017
	EQ 80MG BASE	A208988 004	Aug 22, 2017
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
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ZOLEDRONIC ACID

INJECTABLE; INTRAVENOUS

ZOLEDRONIC ACID

ACTAVIS INC	EQ 4MG BASE/5ML	A202472 001	Mar 04, 2013
DR REDDYS LABS LTD	EQ 4MG BASE/100ML	A204344 001	Nov 19, 2018
SHILPA MEDICARE LTD	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
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ZOLMITRIPTAN

TABLET;ORAL

ZOLMITRIPTAN

ANI PHARMS INC	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
MACLEODS PHARMS LTD	2.5MG	A203772 001	Sep 30, 2015
	5MG	A203772 002	Sep 30, 2015
MYLAN PHARMS INC	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

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
MYLAN	2.5MG	A202855 001	Sep 20, 2019
	5MG	A202855 002	Sep 20, 2019

ZOLPIDEM TARTRATE

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
MYLAN	5MG	A076578 001	Apr 23, 2007

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

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** See List Footnote

ZOLPIDEM TARTRATE

TABLET;ORAL

ZOLPIDEM TARTRATE

	10MG	A076578 002	Apr 23, 2007
MYLAN PHARMS INC	5MG	A078016 001	Apr 23, 2007
	10MG	A078016 002	Apr 23, 2007
STRIDES PHARMA	5MG	A076062 001	Apr 23, 2007
	10MG	A076062 002	Apr 23, 2007
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
	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

TABLET;SUBLINGUAL

INTERMEZZO

+	PURDUE PHARMA	1.75MG	N022328 001	Nov 23, 2011
+		3.5MG	N022328 002	Nov 23, 2011

TABLET, EXTENDED RELEASE;ORAL

ZOLPIDEM TARTRATE

	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

+	SUNOVION PHARMS INC	50MG **	N020789 002	Aug 22, 2003
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ZONISAMIDE

	ANI PHARMS INC	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
AUROBINDO PHARMA LTD	25MG	A077645 002	Sep 29, 2006	
	50MG	A077645 003	Sep 29, 2006	
	100MG	A077645 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	25MG	A077650 001	Apr 20, 2006	
	50MG	A077650 002	Apr 20, 2006	
	100MG	A077650 003	Apr 20, 2006	
MYLAN	25MG	A077637 001	Dec 22, 2005	
	50MG	A077637 002	Dec 22, 2005	
	100MG	A077637 003	Dec 22, 2005	
MYLAN PHARMS INC	25MG	A077647 001	Dec 22, 2005	
	50MG	A077647 002	Dec 22, 2005	
	100MG	A077647 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	

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

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, ARIPIPRAZOLE
 ABILIFY MAINTENA KIT, ARIPIPRAZOLE
 ABILIFY MYCITE KIT, ARIPIPRAZOLE
 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
 ACCUNEB, ALBUTEROL SULFATE
 ACCUPRIL, QUINAPRIL HYDROCHLORIDE
 ACCURETIC, HYDROCHLOROTHIAZIDE
 ACEBUTOLOL HYDROCHLORIDE, ACEBUTOLOL HYDROCHLORIDE
 ACEPHEN, ACETAMINOPHEN (OTC)
 ACETADOTE, ACETYLCYSTEINE
 ACETAMINOPHEN, ACETAMINOPHEN (OTC)
 ACETAMINOPHEN, ACETAMINOPHEN
 ACETAMINOPHEN AND CODEINE PHOSPHATE, ACETAMINOPHEN
 ACETAMINOPHEN, ASPIRIN AND CAFFEINE, ACETAMINOPHEN (OTC)
 ACETAMINOPHEN, CAFFEINE AND DIHYDROCODEINE BITARTRATE, ACETAMINOPHEN
 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
 ACIPHEX SPRINKLE, RABEPRAZOLE SODIUM
 ACITRETIN, ACITRETIN
 ACTHREL, CORTICORELIN OVINE TRIFLUTATE
 ACTICLATE, DOXYCYCLINE HYCLATE
 ACTIGALL, URSODIOL
 ACTIQ, FENTANYL CITRATE
 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
 ADALAT CC, NIFEDIPINE
 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

APPENDIX A - PRODUCT NAME INDEX

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ADEFOVIR DIPIVOXIL, ADEFOVIR DIPIVOXIL
 ADEMPAS, RIOCIQUAT
 ADENOSINE, ADENOSINE
 ADHANSIA XR, METHYLPHENIDATE HYDROCHLORIDE
 ADIPEX-P, PHENTERMINE HYDROCHLORIDE
 ADLYXIN, LIXISENATIDE
 ADMELOG, INSULIN LISPRO
 ADMELOG SOLOSTAR, INSULIN LISPRO
 ADRENACLICK, EPINEPHRINE
 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-SYMPOM COLD & FLU, CHLORPHENIRAMINE MALEATE (OTC)
 ADVIL PM, DIPHENHYDRAMINE CITRATE (OTC)
 ADVIL PM, DIPHENHYDRAMINE HYDROCHLORIDE (OTC)
 ADZENYS ER, AMPHETAMINE
 ADZENYS XR-ODT, AMPHETAMINE
 AEMCOLO, RIFAMYCIN SODIUM
 AFINITOR, EVEROLIMUS
 AFINITOR DISPERZ, EVEROLIMUS
 AFIRMELLE, ETHINYL ESTRADIOL
 AFREZZA, INSULIN RECOMBINANT HUMAN
 AGGRASTAT, TIROFIBAN HYDROCHLORIDE
 AGGRENOLX, ASPIRIN
 AGRYLIN, ANAGRELIDE HYDROCHLORIDE
 AIRDUO DIGIHALER, FLUTICASONE PROPIONATE
 AIRDUO RESPICLICK, FLUTICASONE PROPIONATE
 AK-FLUOR 10%, FLUORESCEIN SODIUM
 AK-FLUOR 25%, FLUORESCEIN SODIUM
 AKBETA, LEVOBUNOLOL HYDROCHLORIDE
 AKLIEF, TRIFAROTENE
 AKOAZ, EPHEDRINE SULFATE
 AKPENTOLATE, CYCLOPENTOLATE HYDROCHLORIDE
 AKTEN, LIDOCAINE HYDROCHLORIDE
 AKTOB, TOBRAMYCIN
 AKYNZEO, FOSNETUPITANT CHLORIDE HYDROCHLORIDE
 AKYNZEO, NETUPITANT
 ALA-CORT, HYDROCORTISONE
 ALA-SCALP, HYDROCORTISONE
 ALAVERT, LORATADINE (OTC)
 ALAWAY, KETOTIFEN FUMARATE (OTC)
 ALBENDAZOLE, ALBENDAZOLE
 ALBENZA, ALBENDAZOLE
 ALBUTEROL SULFATE, ALBUTEROL SULFATE
 ALBUTEROL SULFATE AND IPRATROPIUM BROMIDE, ALBUTEROL SULFATE
 ALCAINE, PROPARACAINE HYDROCHLORIDE
 ALCLOMETASONE DIPROPIONATE, ALCLOMETASONE DIPROPIONATE
 ALDACTAZIDE, HYDROCHLOROTHIAZIDE
 ALDACTONE, SPIRONOLACTONE
 ALDARA, IMIQUIMOD
 ALECENSA, ALECTINIB HYDROCHLORIDE
 ALENDRONATE SODIUM, ALENDRONATE SODIUM
 ALEVE, NAPROXEN SODIUM (OTC)

APPENDIX A - PRODUCT NAME INDEX

** A **

ALEVE PM, DIPHENHYDRAMINE HYDROCHLORIDE (OTC)
 ALEVE-D SINUS & COLD, NAPROXEN SODIUM (OTC)
 ALFENTA, ALFENTANIL HYDROCHLORIDE
 ALFENTANIL, ALFENTANIL HYDROCHLORIDE
 ALFUZOSIN HYDROCHLORIDE, ALFUZOSIN HYDROCHLORIDE
 ALIMTA, PEMETREXED DISODIUM
 ALINIA, NITAZOXANIDE
 ALIQOPA, COPANLISIB DIHYDROCHLORIDE
 ALISKIREN HEMIFUMARATE, ALISKIREN HEMIFUMARATE
 ALKERAN, MELPHALAN
 ALKERAN, MELPHALAN HYDROCHLORIDE
 ALLEGRA ALLERGY, 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
 ALLZITAL, ACETAMINOPHEN
 ALMOTRIPTAN MALATE, ALMOTRIPTAN MALATE
 ALOCRI, NEDOCROMIL SODIUM
 ALOMIDE, LODOXAMIDE TROMETHAMINE
 ALOPRIM, ALLOPURINOL SODIUM
 ALORA, ESTRADIOL
 ALOSETRON HYDROCHLORIDE, ALOSETRON HYDROCHLORIDE
 ALOXI, PALONOSETRON HYDROCHLORIDE
 ALPHAGAN P, BRIMONIDINE TARTRATE
 ALPRAZOLAM, ALPRAZOLAM
 ALPROSTADIL, ALPROSTADIL
 ALREX, LOTEHPREDNOL ETABONATE
 ALTABAX, RETAPAMULIN
 ALTACE, RAMIPRIL
 ALTAFLUOR BENOX, BENOXINATE HYDROCHLORIDE
 ALTAVERA, ETHINYL ESTRADIOL
 ALTOPREV, LOVASTATIN
 ALTRENO, TRETINOIN
 ALUNBRIG, BRIGATINIB
 ALVESCO, CICLESONIDE
 ALVIMOPAN, ALVIMOPAN
 ALYACEN 1/35, ETHINYL ESTRADIOL
 ALYACEN 7/7/7, ETHINYL ESTRADIOL
 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
 AMERGE, NARATRIPTAN HYDROCHLORIDE
 AMICAR, AMINOCAPROIC ACID
 AMIDATE, ETOMIDATE
 AMIFOSTINE, AMIFOSTINE
 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

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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
 AMMONUL, SODIUM BENZOATE
 AMNESTEEM, ISOTRETINOIN
 AMOXAPINE, AMOXAPINE
 AMOXICILLIN, AMOXICILLIN
 AMOXICILLIN AND CLAVULANATE POTASSIUM, AMOXICILLIN
 AMOXICILLIN PEDIATRIC, AMOXICILLIN
 AMOXIL, AMOXICILLIN
 AMPHADASE, HYALURONIDASE
 AMPHETAMINE SULFATE, AMPHETAMINE SULFATE
 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
 AMYVID, FLORBETAPIR F-18
 AMZEEQ, MINOCYCLINE HYDROCHLORIDE
 AN-SULFUR COLLOID, TECHNETIUM TC-99M SULFUR COLLOID KIT
 ANADROL-50, OXYMETHOLONE
 ANAFRANIL, CLOMIPRAMINE HYDROCHLORIDE
 ANAGRELIDE HYDROCHLORIDE, ANAGRELIDE HYDROCHLORIDE
 ANAPROX DS, NAPROXEN SODIUM
 ANASTROZOLE, ANASTROZOLE
 ANCEF IN PLASTIC CONTAINER, CEFAZOLIN SODIUM
 ANCOBON, FLUCYTOSINE
 ANDRODERM, TESTOSTERONE
 ANDROGEL, TESTOSTERONE
 ANDROID 25, METHYLTESTOSTERONE
 ANECTINE, SUCCINYLCHOLINE CHLORIDE
 ANEXSIA 5/325, ACETAMINOPHEN
 ANEXSIA 7.5/325, ACETAMINOPHEN
 ANGELIQ, DROSPIRENONE
 ANGIOMAX, BIVALIRUDIN
 ANGIOMAX RTU, BIVALIRUDIN
 ANJESO, MELOXICAM
 ANNOVERA, ETHINYL ESTRADIOL
 ANORO ELLIPTA, UMECLIDINIUM BROMIDE
 ANTABUSE, DISULFIRAM
 ANTARA (MICRONIZED), FENOFIBRATE
 ANTHELIOS 20, AVOBENZONE (OTC)
 ANTHELIOS 40, AVOBENZONE (OTC)
 ANTHELIOS SX, AVOBENZONE (OTC)
 ANTIVERT, MECLIZINE HYDROCHLORIDE
 ANUSOL HC, HYDROCORTISONE
 APADAZ, ACETAMINOPHEN
 APIDRA, INSULIN GLULISINE RECOMBINANT
 APIDRA SOLOSTAR, INSULIN GLULISINE RECOMBINANT
 APIXABAN, APIXABAN
 APLENZIN, BUPROPION HYDROBROMIDE
 APOKYN, APOMORPHINE HYDROCHLORIDE
 APRACLONIDINE HYDROCHLORIDE, APRACLONIDINE HYDROCHLORIDE

APPENDIX A - PRODUCT NAME INDEX

** A **

APREPITANT, APREPITANT
 APRISO, MESALAMINE
 APTENSIO XR, METHYLPHENIDATE HYDROCHLORIDE
 APTIOM, ESLICARBAZEPINE ACETATE
 APTIVUS, TIPRANAVIR
 AQUASOL A, VITAMIN A PALMITATE
 ARAKODA, TAFENOQUINE SUCCINATE
 ARANELLE, ETHINYL ESTRADIOL
 ARAVA, LEFLUNOMIDE
 ARAZLO, TAZAROTENE
 ARCAPTA NEOHALER, INDACATEROL MALEATE
 ARESTIN, MINOCYCLINE HYDROCHLORIDE
 ARGATROBAN, ARGATROBAN
 ARGATROBAN IN 0.9% SODIUM CHLORIDE, 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
 ARISTOSPAN, TRIAMCINOLONE HEXACETONIDE
 ARIXTRA, FONDAPARINUX SODIUM
 ARMODAFINIL, ARMODAFINIL
 ARNUITY ELLIPTA, FLUTICASONE FUROATE
 AROMASIN, EXEMESTANE
 ARRANON, NELARABINE
 ARSENIC TRIOXIDE, ARSENIC TRIOXIDE
 ARTHROTEC, DICLOFENAC SODIUM
 ASACOL HD, MESALAMINE
 ASCLERA, POLIDOCANOL
 ASCOR, ASCORBIC ACID
 ASHLYNA, ETHINYL ESTRADIOL
 ASMANEX HFA, MOMETASONE FUROATE
 ASMANEX TWISTHALER, MOMETASONE FUROATE
 ASPIRIN AND DIPYRIDAMOLE, ASPIRIN
 ASTAGRAF XL, TACROLIMUS
 ASTELIN, AZELASTINE HYDROCHLORIDE
 ASTEPRO, AZELASTINE HYDROCHLORIDE
 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
 ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM
 ATOVAQUONE, ATOVAQUONE
 ATOVAQUONE AND PROGUANIL HYDROCHLORIDE, ATOVAQUONE
 ATRACURIUM BESYLATE, ATRACURIUM BESYLATE
 ATRACURIUM BESYLATE PRESERVATIVE FREE, ATRACURIUM BESYLATE
 ATRALIN, TRETINOIN
 ATRIDOX, DOXYCYCLINE HYCLATE
 ATRIPLA, EFAVIRENZ
 ATROPEN, ATROPINE
 ATROPINE SULFATE, ATROPINE SULFATE
 ATROPINE SULFATE ANSYR PLASTIC SYRINGE, ATROPINE SULFATE
 ATROPINE SULFATE LIFESHIELD ABOJECT SYRINGE, ATROPINE SULFATE
 ATROVENT HFA, IPRATROPIUM BROMIDE
 AUBAGIO, TERIFLUNOMIDE
 AUGMENTIN '125', AMOXICILLIN
 AUGMENTIN '250', AMOXICILLIN

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** A **

AUGMENTIN '875', AMOXICILLIN
 AUGMENTIN XR, AMOXICILLIN
 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
 AUVI-Q, EPINEPHRINE
 AVAGARD, ALCOHOL (OTC)
 AVAGE, TAZAROTENE
 AVALIDE, HYDROCHLOROTHIAZIDE
 AVANDIA, ROSIGLITAZONE MALEATE
 AVAPRO, IRBESARTAN
 AVC, SULFANILAMIDE
 AVEED, TESTOSTERONE UNDECANOATE
 AVELOX, MOXIFLOXACIN HYDROCHLORIDE
 AVIANE-28, ETHINYL ESTRADIOL
 AVITA, TRETINOIN
 AVODART, DUTASTERIDE
 AVYCAZ, AVIBACTAM SODIUM
 AXID AR, NIZATIDINE (OTC)
 AXUMIN, FLUCICLOVINE F-18
 AYUNA, ETHINYL ESTRADIOL
 AYVAKIT, AVAPRITINIB
 AZACITIDINE, AZACITIDINE
 AZACTAM, AZTREONAM
 AZASAN, AZATHIOPRINE
 AZASITE, AZITHROMYCIN
 AZATHIOPRINE, AZATHIOPRINE
 AZATHIOPRINE SODIUM, AZATHIOPRINE SODIUM
 AZEDRA, IOBENGUANE I-131
 AZELAIC ACID, AZELAIC ACID
 AZELASTINE HYDROCHLORIDE, AZELASTINE HYDROCHLORIDE
 AZELASTINE HYDROCHLORIDE AND FLUTICASONE PROPIONATE, AZELASTINE HYDROCHLORIDE
 AZELEX, AZELAIC ACID
 AZILECT, RASAGILINE MESYLATE
 AZITHROMYCIN, AZITHROMYCIN
 AZOPT, BRINZOLAMIDE
 AZOR, AMLODIPINE BESYLATE
 AZTREONAM, AZTREONAM
 AZULFIDINE, SULFASALAZINE
 AZULFIDINE EN-TABS, SULFASALAZINE

** B **

BACIIM, BACITRACIN
 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
 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

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** B **

BARHEMSYS, AMISULPRIDE
 BASAGLAR, INSULIN GLARGINE
 BAXDELA, DELAFLOXACIN MEGLUMINE
 BECONASE AQ, BECLOMETHASONE DIPROPIONATE MONOHYDRATE
 BEKYREE, DESOGESTREL
 BELBUCA, BUPRENORPHINE HYDROCHLORIDE
 BELEODAQ, BELINOSTAT
 BELRAPZO, BENDAMUSTINE HYDROCHLORIDE
 BELSOMRA, SUVOREXANT
 BENAZEPRIL HYDROCHLORIDE, BENAZEPRIL HYDROCHLORIDE
 BENAZEPRIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, BENAZEPRIL HYDROCHLORIDE
 BENDEKA, BENDAMUSTINE HYDROCHLORIDE
 BENICAR, OLMESARTAN MEDOXOMIL
 BENICAR HCT, HYDROCHLOROTHIAZIDE
 BENTYL, DICYCLOMINE HYDROCHLORIDE
 BENTYL PRESERVATIVE FREE, DICYCLOMINE HYDROCHLORIDE
 BENZACLIN, BENZOYL PEROXIDE
 BENZAMYCIN, BENZOYL PEROXIDE
 BENZNIDAZOLE, BENZNIDAZOLE
 BENZONATATE, BENZONATATE
 BENZPHETAMINE HYDROCHLORIDE, BENZPHETAMINE HYDROCHLORIDE
 BENZTROPINE MESYLATE, BENZTROPINE MESYLATE
 BEPREVE, BEPOTASTINE BESILATE
 BESIVANCE, BESIFLOXACIN HYDROCHLORIDE
 BETA-VAL, BETAMETHASONE VALERATE
 BETADINE, POVIDONE-IODINE
 BETAGAN, LEVOBUNOLOL HYDROCHLORIDE
 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
 BEVYXXA, BETRIXABAN
 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
 BILTRICIDE, PRAZIQUANTEL
 BIMATOPROST, BIMATOPROST
 BINOSTO, ALENDRONATE SODIUM
 BIORPHEN, PHENYLEPHRINE HYDROCHLORIDE
 BIOSCRUB, CHLORHEXIDINE GLUCONATE (OTC)
 BISMUTH SUBSALICYLATE, METRONIDAZOLE AND TETRACYCLINE HYDROCHLORIDE, BISMUTH SUBSALICYLATE
 BISOPROLOL FUMARATE, BISOPROLOL FUMARATE
 BISOPROLOL FUMARATE AND HYDROCHLOROTHIAZIDE, BISOPROLOL FUMARATE
 BIVALIRUDIN, BIVALIRUDIN
 BIVALIRUDIN IN 0.9% SODIUM CHLORIDE, BIVALIRUDIN
 BLEOMYCIN SULFATE, BLEOMYCIN SULFATE
 BLEPH-10, SULFACETAMIDE SODIUM
 BLEPHAMIDE, PREDNISOLONE ACETATE
 BLEPHAMIDE S.O.P., PREDNISOLONE ACETATE
 BLISOVI 24 FE, ETHINYL ESTRADIOL

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BLISOVI FE 1.5/30, ETHINYL ESTRADIOL
 BLISOVI FE 1/20, ETHINYL ESTRADIOL
 BLOXIVERZ, NEOSTIGMINE METHYLSULFATE
 BONIVA, IBANDRONATE SODIUM
 BONJESTA, DOXYLAMINE SUCCINATE
 BONSITY, TERIPARATIDE
 BONTRIL PDM, PHENDIMETRAZINE TARTRATE
 BORTEZOMIB, BORTEZOMIB
 BOSENTAN, BOSENTAN
 BOSULIF, BOSUTINIB MONOHYDRATE
 BRAFTOVI, ENCORAFENIB
 BREO ELLIPTA, FLUTICASON FUROATE
 BRETHINE, TERBUTALINE SULFATE
 BRETILIUM TOSYLATE, BRETILIUM TOSYLATE
 BREVIBLOC, ESMOLOL HYDROCHLORIDE
 BREVIBLOC DOUBLE STRENGTH IN PLASTIC CONTAINER, ESMOLOL HYDROCHLORIDE
 BREVIBLOC IN PLASTIC CONTAINER, ESMOLOL HYDROCHLORIDE
 BREVICON 28-DAY, ETHINYL ESTRADIOL
 BREVITAL SODIUM, METHOHEXITAL SODIUM
 BRIAN CARE, CHLORHEXIDINE GLUCONATE (OTC)
 BRIDION, SUGAMMADEX SODIUM
 BRIELLYN, ETHINYL ESTRADIOL
 BRILINTA, TICAGRELOR
 BRIMONIDINE TARTRATE, BRIMONIDINE TARTRATE
 BRISDELLE, PAROXETINE MESYLATE
 BRIVIACT, BRIVARACETAM
 BROMFED-DM, BROMPHENIRAMINE MALEATE
 BROMFENAC SODIUM, BROMFENAC SODIUM
 BROMOCRIPTINE MESYLATE, BROMOCRIPTINE MESYLATE
 BROMPHENIRAMINE MALEATE, PSEUDOEPHEDRINE HYDROCHLORIDE AND DEXTROMETHORPHAN HYDROBROMIDE,
 BROMSITE, BROMFENAC SODIUM
 BRONCHO SALINE, SODIUM CHLORIDE (OTC)
 BROVANA, ARFORMOTEROL TARTRATE
 BRUKINSA, ZANUBRUTINIB
 BRYHALI, HALOBETASOL PROPIONATE
 BSS, CALCIUM CHLORIDE
 BSS PLUS, CALCIUM CHLORIDE
 BUDESONIDE, BUDESONIDE (OTC)
 BUDESONIDE, BUDESONIDE
 BUMETANIDE, BUMETANIDE
 BUMEX, BUMETANIDE
 BUNAVAIL, BUPRENORPHINE HYDROCHLORIDE
 BUPHENYL, SODIUM PHENYLBUTYRATE
 BUPIVACAINE HYDROCHLORIDE, BUPIVACAINE HYDROCHLORIDE
 BUPIVACAINE HYDROCHLORIDE AND EPINEPHRINE, BUPIVACAINE HYDROCHLORIDE
 BUPIVACAINE HYDROCHLORIDE PRESERVATIVE FREE, BUPIVACAINE HYDROCHLORIDE
 BUPRENEX, BUPRENORPHINE HYDROCHLORIDE
 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)
 BUTISOL SODIUM, BUTABARBITAL SODIUM
 BUTORPHANOL TARTRATE, BUTORPHANOL TARTRATE
 BUTORPHANOL TARTRATE PRESERVATIVE FREE, BUTORPHANOL TARTRATE
 BUTRANS, BUPRENORPHINE

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BYDUREON, EXENATIDE SYNTHETIC
 BYDUREON BCISE, EXENATIDE
 BYDUREON PEN, EXENATIDE SYNTHETIC
 BYETTA, EXENATIDE SYNTHETIC
 BYNFEZIA PEN, OCTREOTIDE ACETATE
 BYSTOLIC, NEBIVOLOL HYDROCHLORIDE

** C **

CABERGOLINE, CABERGOLINE
 CABOMETYX, CABOZANTINIB S-MALATE
 CADUET, AMLODIPINE BESYLATE
 CAFKIT, CAFFEINE CITRATE
 CAFERGOT, CAFFEINE
 CAFFEINE CITRATE, CAFFEINE CITRATE
 CALAN SR, VERAPAMIL HYDROCHLORIDE
 CALCIPOTRIENE, CALCIPOTRIENE
 CALCIPOTRIENE AND BETAMETHASONE 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
 CALCIUM DISODIUM VERSENATE, EDETATE CALCIUM DISODIUM
 CALCIUM GLUCONATE, CALCIUM GLUCONATE
 CALCIUM GLUCONATE IN SODIUM CHLORIDE, CALCIUM GLUCONATE
 CALDOLOR, IBUPROFEN
 CALQUENCE, ACALABRUTINIB
 CAMBIA, DICLOFENAC POTASSIUM
 CAMILA, NORETHINDRONE
 CAMPTOSAR, IRINOTECAN HYDROCHLORIDE
 CANASA, MESALAMINE
 CANCIDAS, CASPOFUNGIN ACETATE
 CANDESARTAN CILEXETIL, CANDESARTAN CILEXETIL
 CANDESARTAN CILEXETIL AND HYDROCHLOROTHIAZIDE, CANDESARTAN CILEXETIL
 CAPASTAT SULFATE, CAPREOMYCIN SULFATE
 CAPECITABINE, CAPECITABINE
 CAPEX, FLUOCINOLONE ACETONIDE
 CAPITAL SOLEIL 15, AVOBENZONE (OTC)
 CAPLYTA, LUMATEPERONE TOSYLATE
 CAPRELSA, VANDETANIB
 CAPREOMYCIN SULFATE, CAPREOMYCIN SULFATE
 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

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CARDURA XL, DOXAZOSIN MESYLATE
 CARFILZOMIB, CARFILZOMIB
 CARISOPRODOL, CARISOPRODOL
 CARISOPRODOL AND ASPIRIN, ASPIRIN
 CARISOPRODOL, ASPIRIN AND CODEINE PHOSPHATE, ASPIRIN
 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
 CATAPRES, CLONIDINE HYDROCHLORIDE
 CATAPRES-TTS-1, CLONIDINE
 CATAPRES-TTS-2, CLONIDINE
 CATAPRES-TTS-3, CLONIDINE
 CAVERJECT, ALPROSTADIL
 CAVERJECT IMPULSE, ALPROSTADIL
 CAYSTON, AZTREONAM
 CEFACLOR, CEFACLOR
 CEFADROXIL, CEFADROXIL/CEFADROXIL HEMIHYDRATE
 CEFAZOLIN AND DEXTROSE, CEFAZOLIN SODIUM
 CEFAZOLIN IN PLASTIC CONTAINER, 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
 CEFOTAXIME, CEFOTAXIME SODIUM
 CEFOTETAN, CEFOTETAN DISODIUM
 CEFOTETAN AND DEXTROSE IN DUPLEX CONTAINER, 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
 CEFTAZIDIME IN DEXTROSE CONTAINER, CEFTAZIDIME
 CEFTRIAXONE, CEFTRIAXONE SODIUM
 CEFTRIAXONE AND DEXTROSE IN DUPLEX CONTAINER, CEFTRIAXONE SODIUM
 CEFTRIAXONE IN PLASTIC CONTAINER, CEFTRIAXONE SODIUM
 CEFTRIAXONE SODIUM, CEFTRIAXONE SODIUM
 CEFUROXIME AND DEXTROSE IN DUPLEX CONTAINER, CEFUROXIME SODIUM
 CEFUROXIME AXETIL, CEFUROXIME AXETIL
 CEFUROXIME SODIUM, CEFUROXIME SODIUM
 CELEBREX, 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

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CEREZYME, IMIGLUCERASE
 CERINTA, ETHINYL ESTRADIOL
 CERUBIDINE, DAUNORUBICIN HYDROCHLORIDE
 CERVIDIL, DINOPROSTONE
 CESAMET, NABILONE
 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
 CETROTIDE, CETRORELIX
 CEVIMELINE HYDROCHLORIDE, CEVIMELINE HYDROCHLORIDE
 CHANTIX, VARENICLINE TARTRATE
 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 ALLEGRA ALLERGY, FEXOFENADINE 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)
 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 RELIEF, CETIRIZINE HYDROCHLORIDE (OTC)
 CHIRHOSTIM, SECRETIN SYNTHETIC HUMAN
 CHLORAMPHENICOL SODIUM SUCCINATE, CHLORAMPHENICOL SODIUM SUCCINATE
 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
 CHLORHEXIDINE GLUCONATE, CHLORHEXIDINE GLUCONATE (OTC)
 CHLORHEXIDINE GLUCONATE, CHLORHEXIDINE GLUCONATE
 CHLOROPROCAINE HYDROCHLORIDE, CHLOROPROCAINE HYDROCHLORIDE
 CHLOROQUINE PHOSPHATE, CHLOROQUINE PHOSPHATE
 CHLOROTHIAZIDE, CHLOROTHIAZIDE
 CHLOROTHIAZIDE SODIUM, CHLOROTHIAZIDE SODIUM
 CHLORPHENIRAMINE MALEATE, CHLORPHENIRAMINE MALEATE (OTC)
 CHLORPROMAZINE HYDROCHLORIDE, CHLORPROMAZINE HYDROCHLORIDE
 CHLORTHALIDONE, CHLORTHALIDONE
 CHLORZOAZONE, CHLORZOAZONE
 CHOLAC, LACTULOSE
 CHOLBAM, CHOLIC ACID
 CHOLESTYRAMINE, CHOLESTYRAMINE
 CHOLESTYRAMINE LIGHT, CHOLESTYRAMINE
 CHOLETEC, TECHNETIUM TC-99M MEBROFENIN KIT
 CHOLINE C-11, CHOLINE C-11
 CHORIONIC GONADOTROPIN, GONADOTROPIN, CHORIONIC
 CHROMIC CHLORIDE IN PLASTIC CONTAINER, CHROMIC CHLORIDE
 CIALIS, TADALAFIL
 CICLOPIROX, CICLOPIROX
 CIDA-STAT, CHLORHEXIDINE GLUCONATE (OTC)
 CIDOFOVIR, CIDOFOVIR
 CILOSTAZOL, CILOSTAZOL

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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 EXTENDED RELEASE, CIPROFLOXACIN
 CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE
 CIPROFLOXACIN IN DEXTROSE 5% IN PLASTIC CONTAINER, 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 24 HOUR, DESLORATADINE
 CLARINEX-D 12 HOUR, DESLORATADINE
 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 (OTC)
 CLEMASTINE FUMARATE, CLEMASTINE FUMARATE
 CLENPIQ, CITRIC ACID
 CLEOCIN, CLINDAMYCIN PALMITATE HYDROCHLORIDE
 CLEOCIN, CLINDAMYCIN PHOSPHATE
 CLEOCIN HYDROCHLORIDE, CLINDAMYCIN HYDROCHLORIDE
 CLEOCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
 CLEOCIN PHOSPHATE IN DEXTROSE 5% IN PLASTIC CONTAINER, 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

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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 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
 CLODERM, CLOCORTOLONE PIVALATE
 CLOFARABINE, CLOFARABINE
 CLOLAR, CLOFARABINE
 CLOMIPHENE CITRATE, CLOMIPHENE CITRATE
 CLOMIPRAMINE HYDROCHLORIDE, CLOMIPRAMINE HYDROCHLORIDE
 CLONAZEPAM, CLONAZEPAM
 CLONIDINE, CLONIDINE
 CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
 CLOPIDOGREL BISULFATE, CLOPIDOGREL BISULFATE
 CLORAZEPATE DIPOTASSIUM, CLORAZEPATE DIPOTASSIUM
 CLOROTEKAL, CHLOROPROCAINE HYDROCHLORIDE
 CLOTRIMAZOLE, CLOTRIMAZOLE (OTC)
 CLOTRIMAZOLE, CLOTRIMAZOLE
 CLOTRIMAZOLE AND BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
 CLOVIQUE, TRIENTINE HYDROCHLORIDE
 CLOZAPINE, CLOZAPINE
 CLOZARIL, CLOZAPINE
 COARTEM, ARTEMETHER
 CODEINE SULFATE, CODEINE SULFATE
 COGENTIN, BENZTROPINE MESYLATE
 COL-PROBENECID, COLCHICINE
 COLAZAL, BALSALAZIDE DISODIUM
 COLCHICINE, COLCHICINE
 COLCRYS, 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
 COLYTE WITH FLAVOR PACKS, POLYETHYLENE GLYCOL 3350
 COMBIGAN, BRIMONIDINE TARTRATE
 COMBIPATCH, ESTRADIOL
 COMBIVENT RESPIMAT, ALBUTEROL SULFATE
 COMBIVIR, LAMIVUDINE
 COMETRIQ, CABOZANTINIB S-MALATE
 COMPLERA, EMTRICITABINE
 COMPRO, PROCHLORPERAZINE
 COMTAN, ENTACAPONE
 CONCERTA, METHYLPHENIDATE HYDROCHLORIDE

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CONDYLOX, PODOFILOX
 CONJUPRI, LEVAMLODIPINE MALEATE
 CONRAY, IOTHALAMATE MEGLUMINE
 CONSENSI, AMLODIPINE BESYLATE
 CONSTILAC, LACTULOSE
 CONTRAVE, BUPROPION HYDROCHLORIDE
 CONZIP, TRAMADOL HYDROCHLORIDE
 COPAXONE, GLATIRAMER ACETATE
 COPIKTRA, DUVELISIB
 CORDRAN, FLURANDRENOLIDE
 CORDRAN SP, FLURANDRENOLIDE
 COREG, CARVEDILOL
 COREG CR, CARVEDILOL PHOSPHATE
 CORGARD, NADOLOL
 CORLANOR, IVABRADINE
 CORLANOR, IVABRADINE HYDROCHLORIDE
 CORLOPAM, FENOLDOPAM MESYLATE
 CORMAX, CLOBETASOL PROPIONATE
 CORPHEDRA, EPHEDRINE SULFATE
 CORTEF, HYDROCORTISONE
 CORTENEMA, HYDROCORTISONE
 CORTIFOAM, HYDROCORTISONE ACETATE
 CORTISONE ACETATE, CORTISONE ACETATE
 CORTISPORIN, BACITRACIN ZINC
 CORTISPORIN, HYDROCORTISONE ACETATE
 CORTROSYN, COSYNTROPIN
 CORVERT, IBUTILIDE FUMARATE
 CORZIDE, BENDROFLUMETHIAZIDE
 COSMEGEN, DACTINOMYCIN
 COSOPT, DORZOLAMIDE HYDROCHLORIDE
 COSOPT PF, DORZOLAMIDE HYDROCHLORIDE
 COSYNTROPIN, COSYNTROPIN
 COTELLIC, COBIMETINIB FUMARATE
 COTEMPLA XR-ODT, METHYLPHENIDATE
 COUMADIN, WARFARIN SODIUM
 COZAAR, LOSARTAN POTASSIUM
 CREON, PANCRELIPASE (AMYLASE
 CRESEMBA, ISAVUCONAZONIUM SULFATE
 CRESTOR, ROSUVASTATIN CALCIUM
 CRINONE, PROGESTERONE
 CRIXIVAN, INDINAVIR SULFATE
 CROMOLYN SODIUM, CROMOLYN SODIUM (OTC)
 CROMOLYN SODIUM, CROMOLYN SODIUM
 CROTAN, CROTAMITON
 CRYSELLE, ETHINYL ESTRADIOL
 CUBICIN, DAPTOMYCIN
 CUBICIN RF, DAPTOMYCIN
 CUPRIC CHLORIDE IN PLASTIC CONTAINER, CUPRIC CHLORIDE
 CUPRIMINE, PENICILLAMINE
 CUROSURF, PORACTANT ALFA
 CUTIVATE, FLUTICASONE PROPIONATE
 CUVPOSA, GLYCOPYRROLATE
 CYANOCOBALAMIN, CYANOCOBALAMIN
 CYANOKIT, HYDROXOCOBALAMIN
 CYCLAFEM 0.5/35, ETHINYL ESTRADIOL
 CYCLAFEM 1/35, ETHINYL ESTRADIOL
 CYCLAFEM 7/7/7, ETHINYL ESTRADIOL
 CYCLESSA, DESOGESTREL
 CYCLOBENZAPRINE HYDROCHLORIDE, CYCLOBENZAPRINE HYDROCHLORIDE
 CYCLOGYL, CYCLOPENTOLATE HYDROCHLORIDE
 CYCLOMYDRIL, CYCLOPENTOLATE HYDROCHLORIDE
 CYCLOPENTOLATE HYDROCHLORIDE, CYCLOPENTOLATE HYDROCHLORIDE
 CYCLOPHOSPHAMIDE, CYCLOPHOSPHAMIDE
 CYCLOSET, BROMOCRIPTINE MESYLATE
 CYCLOSPORINE, CYCLOSPORINE

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CYKLOKAPRON, TRANEXAMIC ACID
 CYMBALTA, DULOXETINE HYDROCHLORIDE
 CYONANZ, ETHINYL ESTRADIOL
 CYPROHEPTADINE HYDROCHLORIDE, CYPROHEPTADINE HYDROCHLORIDE
 CYSTADANE, BETAINE
 CYSTAGON, CYSTEAMINE BITARTRATE
 CYSTARAN, CYSTEAMINE HYDROCHLORIDE
 CYSTO-CONRAY II, IOTHALAMATE MEGLUMINE
 CYSTOGRAFIN, DIATRIZOATE MEGLUMINE
 CYSTOGRAFIN DILUTE, DIATRIZOATE MEGLUMINE
 CYSVIEW KIT, HEXAMINOLEVULINATE HYDROCHLORIDE
 CYTARABINE, CYTARABINE
 CYTOMEL, LIOTHYRONINE SODIUM
 CYTOTEC, MISOPROSTOL
 CYTOVENE, GANCICLOVIR SODIUM

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D.H.E. 45, DIHYDROERGOTAMINE MESYLATE
 DABIGATRAN ETEXILATE MESYLATE, DABIGATRAN ETEXILATE MESYLATE
 DACARBAZINE, DACARBAZINE
 DACOGEN, DECITABINE
 DACTINOMYCIN, DACTINOMYCIN
 DALFAMPRIDINE, DALFAMPRIDINE
 DALIRESP, ROFLUMILAST
 DALVANCE, DALBAVANCIN HYDROCHLORIDE
 DANAZOL, DANAZOL
 DANTRIUM, DANTROLENE SODIUM
 DANTROLENE SODIUM, DANTROLENE SODIUM
 DAPSONE, DAPSONE
 DAPTOMYCIN, DAPTOMYCIN
 DARAPRIM, PYRIMETHAMINE
 DARIFENACIN, DARIFENACIN HYDROBROMIDE
 DARIFENACIN HYDROBROMIDE, DARIFENACIN HYDROBROMIDE
 DARUNAVIR, DARUNAVIR
 DASETTA 1/35, ETHINYL ESTRADIOL
 DASETTA 7/7/7, ETHINYL ESTRADIOL
 DATSCAN, IOFLUPANE I-123
 DAUNORUBICIN HYDROCHLORIDE, DAUNORUBICIN HYDROCHLORIDE
 DAURISMO, GLASDEGIB MALEATE
 DAYPRO, OXAPROZIN
 DAYSEE, ETHINYL ESTRADIOL
 DAYTRANA, METHYLPHENIDATE
 DDAVP, DESMOPRESSIN ACETATE
 DDAVP (NEEDS NO REFRIGERATION), DESMOPRESSIN ACETATE
 DECITABINE, DECITABINE
 DEFERASIROX, DEFERASIROX
 DEFEROXAMINE MESYLATE, DEFEROXAMINE MESYLATE
 DEFINITY, PERFLUTREN
 DEFITELIO, DEFIBROTIDE SODIUM
 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)
 DELZICOL, MESALAMINE
 DEMADAX, TORSEMIDE
 DEMECLOCYCLINE HYDROCHLORIDE, DEMECLOCYCLINE HYDROCHLORIDE
 DEMEROL, MEPERIDINE HYDROCHLORIDE

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DEMSER, METYROSINE
 DENAVIR, PENCICLOVIR
 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
 DERMATOP E EMOLLIENT, PREDNICARBATE
 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
 DESONATE, DESONIDE
 DESONIDE, DESONIDE
 DESOWEN, DESONIDE
 DESOXIMETASONE, DESOXIMETASONE
 DESOXYN, METHAMPHETAMINE HYDROCHLORIDE
 DESVENLAFAXINE, DESVENLAFAXINE
 DESVENLAFAXINE SUCCINATE, DESVENLAFAXINE SUCCINATE
 DETROL, TOLTERODINE TARTRATE
 DETROL LA, TOLTERODINE TARTRATE
 DEXAMETHASONE, DEXAMETHASONE
 DEXAMETHASONE INTENSOL, DEXAMETHASONE
 DEXAMETHASONE SODIUM PHOSPHATE, DEXAMETHASONE SODIUM PHOSPHATE
 DEXAMETHASONE SODIUM PHOSPHATE PRESERVATIVE FREE, DEXAMETHASONE SODIUM PHOSPHATE
 DEXASPORIN, DEXAMETHASONE
 DEXCHLORPHENIRAMINE MALEATE, DEXCHLORPHENIRAMINE MALEATE
 DEXEDRINE, DEXTROAMPHETAMINE SULFATE
 DEXILANT, DEXLANSOPRAZOLE
 DEXLANSOPRAZOLE, DEXLANSOPRAZOLE
 DEXMEDETOMIDINE, DEXMEDETOMIDINE HYDROCHLORIDE
 DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE
 DEXMETHYLPHENIDATE HYDROCHLORIDE, DEXMETHYLPHENIDATE HYDROCHLORIDE
 DEXRAZOXANE HYDROCHLORIDE, DEXRAZOXANE HYDROCHLORIDE
 DEXTENZA, DEXAMETHASONE
 DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE
 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

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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% 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% 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% 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% 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 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
DEXTROSE 50% IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 70% IN PLASTIC CONTAINER, DEXTROSE
DEXYCU KIT, DEXAMETHASONE
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
DIAZEPAM, DIAZEPAM
DIAZEPAM INTENSOL, DIAZEPAM
DIAZOXIDE, DIAZOXIDE
DICLEGIS, DOXYLAMINE SUCCINATE
DICLOFENAC POTASSIUM, DICLOFENAC POTASSIUM
DICLOFENAC SODIUM, DICLOFENAC SODIUM
DICLOFENAC SODIUM AND MISOPROSTOL, DICLOFENAC SODIUM
DICLOXACILLIN SODIUM, DICLOXACILLIN SODIUM
DICYCLOMINE HYDROCHLORIDE, DICYCLOMINE HYDROCHLORIDE
DICYCLOMINE HYDROCHLORIDE (PRESERVATIVE FREE), DICYCLOMINE HYDROCHLORIDE

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DIDANOSINE, DIDANOSINE
DIETHYLPROPION HYDROCHLORIDE, DIETHYLPROPION HYDROCHLORIDE
DIFFERIN, ADAPALENE (OTC)
DIFFERIN, ADAPALENE
DIFICID, FIDAXOMICIN
DIFLORASONE DIACETATE, DIFLORASONE DIACETATE
DIFLUCAN, FLUCONAZOLE
DIFLUNISAL, DIFLUNISAL
DIGOXIN, DIGOXIN
DIHYDROERGOTAMINE MESYLATE, DIHYDROERGOTAMINE MESYLATE
DILANTIN, PHENYTOIN
DILANTIN, PHENYTOIN SODIUM
DILANTIN-125, PHENYTOIN
DILATRATE-SR, ISOSORBIDE DINITRATE
DILAUDID, HYDROMORPHONE HYDROCHLORIDE
DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
DILTZAC, DILTIAZEM HYDROCHLORIDE
DIMENHYDRINATE, DIMENHYDRINATE
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
DITROPAN XL, OXYBUTYNIN CHLORIDE
DIURIL, CHLOROTHIAZIDE
DIURIL, CHLOROTHIAZIDE SODIUM
DIVALPROEX SODIUM, DIVALPROEX SODIUM
DIVIGEL, ESTRADIOL
DOBUTAMINE HYDROCHLORIDE, DOBUTAMINE HYDROCHLORIDE
DOBUTAMINE HYDROCHLORIDE IN DEXTROSE 5% IN PLASTIC CONTAINER, DOBUTAMINE HYDROCHLORIDE
DOCETAXEL, DOCETAXEL
DOCOSANOL, DOCOSANOL (OTC)
DOFETILDE, DOFETILIDE
DOFETILIDE, DOFETILIDE
DOLOPHINE HYDROCHLORIDE, METHADONE HYDROCHLORIDE
DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
DOPAMINE HYDROCHLORIDE, DOPAMINE HYDROCHLORIDE
DOPAMINE HYDROCHLORIDE AND DEXTROSE 5%, DOPAMINE HYDROCHLORIDE
DOPAMINE HYDROCHLORIDE AND DEXTROSE 5% IN PLASTIC CONTAINER, 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

APPENDIX A - PRODUCT NAME INDEX

** D **

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
 DROPERIDOL, DROPERIDOL
 DROSPIRENONE AND ETHINYL ESTRADIOL, DROSPIRENONE
 DROSPIRENONE, ETHINYL ESTRADIOL AND LEVOMEFOLATE CALCIUM, DROSPIRENONE
 DROXIA, HYDROXYUREA
 DSUVIA, SUFENTANIL CITRATE
 DUAC, BENZOYL PEROXIDE
 DUAKLIR PRESSAIR, ACLIDINIUM BROMIDE
 DUAVEE, BAZEDOXIFENE ACETATE
 DUETACT, GLIMEPIRIDE
 DUEXIS, FAMOTIDINE
 DULERA, FORMOTEROL FUMARATE
 DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE
 DUOBRII, HALOBETASOL PROPIONATE
 DUODOTE, ATROPINE
 DUOPA, CARBIDOPA
 DURACLON, CLONIDINE HYDROCHLORIDE
 DURAGESIC-100, FENTANYL
 DURAGESIC-12, FENTANYL
 DURAGESIC-25, FENTANYL
 DURAGESIC-37, FENTANYL
 DURAGESIC-50, FENTANYL
 DURAGESIC-75, FENTANYL
 DURAMORPH PF, MORPHINE SULFATE
 DURAPREP, IODINE POVACRYLEX (OTC)
 DUREZOL, DIFLUPREDNATE
 DURLAZA, ASPIRIN
 DUTASTERIDE, DUTASTERIDE
 DUTASTERIDE AND TAMSULOSIN HYDROCHLORIDE, DUTASTERIDE
 DUTOPROL, HYDROCHLOROTHIAZIDE
 DUVOID, BETHANECHOL CHLORIDE
 DYANAVEL XR, AMPHETAMINE
 DYAZIDE, HYDROCHLOROTHIAZIDE
 DYCLOPRO, DYCLONINE HYDROCHLORIDE
 DYMISTA, AZELASTINE HYDROCHLORIDE
 DYNACIN, MINOCYCLINE HYDROCHLORIDE
 DYRENIUM, TRIAMTERENE

** E **

E-Z SCRUB 201, POVIDONE-IODINE (OTC)
 E-Z SCRUB 241, POVIDONE-IODINE (OTC)
 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
 EDARBI, AZILSARTAN KAMEDOXOMIL
 EDARBYCLOR, AZILSARTAN KAMEDOXOMIL
 EDECRIN, ETHACRYNATE SODIUM
 EDECRIN, ETHACRYNIC ACID
 EDEX, ALPROSTADIL
 EDLUAR, ZOLPIDEM TARTRATE
 EDURANT, RILPIVIRINE HYDROCHLORIDE

APPENDIX A - PRODUCT NAME INDEX

** E **

EFAVIRENZ, EFAVIRENZ
 EFAVIRENZ, EMTRICITABINE, AND TENOFOVIR DISOPROXIL FUMARATE, EFAVIRENZ
 EFFEXOR XR, VENLAFAXINE HYDROCHLORIDE
 EFFIENT, PRASUGREL HYDROCHLORIDE
 EFUDEX, FLUOROURACIL
 EGATEN, TRICLABENDAZOLE
 EGRIFTA, TESAMORELIN ACETATE
 ELCYS, CYSTEINE HYDROCHLORIDE
 ELELYSO, TALIGLUCERASE ALFA
 ELEPSIA XR, LEVETIRACETAM
 ELESTAT, EPINASTINE HYDROCHLORIDE
 ELESTRIN, ESTRADIOL
 ELETRIPTAN HYDROBROMIDE, ELETRIPTAN HYDROBROMIDE
 ELIDEL, PIMECROLIMUS
 ELIFEMME, ETHINYL ESTRADIOL
 ELIGARD, LEUPROLIDE ACETATE
 ELINEST, ETHINYL ESTRADIOL
 ELIQUIS, APIXABAN
 ELIXOPHYLLIN, THEOPHYLLINE
 ELLA, ULIPRISTAL ACETATE
 ELLENCE, EPIRUBICIN HYDROCHLORIDE
 ELLIOTTS B SOLUTION, CALCIUM CHLORIDE
 ELMIRON, PENTOSAN POLYSULFATE SODIUM
 ELOCON, MOMETASONE FUROATE
 ELOXATIN, OXALIPLATIN
 ELURYNG, ETHINYL ESTRADIOL
 EMBELINE, CLOBETASOL PROPIONATE
 EMBELINE E, CLOBETASOL PROPIONATE
 EMCYT, ESTRAMUSTINE PHOSPHATE SODIUM
 EMEND, APREPITANT
 EMEND, FOSAPREPITANT DIMEGLUMINE
 EMFLAZA, DEFLAZACORT
 EMLA, LIDOCAINE
 EMOQUETTE, DESOGESTREL
 EMSAM, SELEGILINE
 EMTRICITABINE, EMTRICITABINE
 EMTRICITABINE AND TENOFOVIR DISOPROXIL FUMARATE, EMTRICITABINE
 EMTRIVA, EMTRICITABINE
 EMVERM, MEBENDAZOLE
 ENABLEX, DARIFENACIN HYDROBROMIDE
 ENALAPRIL MALEATE, ENALAPRIL MALEATE
 ENALAPRIL MALEATE AND HYDROCHLOROTHIAZIDE, ENALAPRIL MALEATE
 ENALAPRILAT, ENALAPRILAT
 ENDARI, L-GLUTAMINE
 ENDOMETRIN, PROGESTERONE
 ENOXAPARIN SODIUM, ENOXAPARIN SODIUM
 ENOXAPARIN SODIUM (PRESERVATIVE FREE), ENOXAPARIN SODIUM
 ENPRESSE-28, ETHINYL ESTRADIOL
 ENSKYCE, DESOGESTREL
 ENSTILAR, BETAMETHASONE DIPROPIONATE
 ENTACAPONE, ENTACAPONE
 ENTECAVIR, ENTECAVIR
 ENTEREG, ALVIMOPAN
 ENTOCORT EC, BUDESONIDE
 ENTRESTO, SACUBITRIL
 ENULOSE, LACTULOSE
 ENVARUS XR, TACROLIMUS
 EOVIIST, GADOXETATE DISODIUM
 EPANED, ENALAPRIL MALEATE
 EPCLUSA, SOFOSBUVIR
 EPHEDRINE SULFATE, EPHEDRINE SULFATE
 EPIDIOLEX, CANNABIDIOL
 EPIDUO, ADAPALENE
 EPIDUO FORTE, ADAPALENE
 EPIFOAM, HYDROCORTISONE ACETATE

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** E **

EPINASTINE HYDROCHLORIDE, EPINASTINE HYDROCHLORIDE
 EPINEPHRINE, EPINEPHRINE
 EPINEPHRINE (AUTOINJECTOR), EPINEPHRINE
 EPINEPHRINE (COPACKAGED), EPINEPHRINE
 EPIPEN, EPINEPHRINE
 EPIPEN JR., EPINEPHRINE
 EPIRUBICIN HYDROCHLORIDE, EPIRUBICIN HYDROCHLORIDE
 EPITOL, CARBAMAZEPINE
 EPIVIR, LAMIVUDINE
 EPIVIR-HBV, LAMIVUDINE
 EPLERENONE, EPLERENONE
 EPOPROSTENOL SODIUM, EPOPROSTENOL SODIUM
 EPROSARTAN MESYLATE, EPROSARTAN MESYLATE
 EPTIFIBATIDE, EPTIFIBATIDE
 EPZICOM, ABACAVIR SULFATE
 EQUETRO, CARBAMAZEPINE
 ERAXIS, ANIDULAFUNGIN
 ERGOCALCIFEROL, ERGOCALCIFEROL
 ERGOLOID MESYLATES, ERGOLOID MESYLATES
 ERGOMAR, ERGOTAMINE TARTRATE
 ERGOTAMINE TARTRATE AND CAFFEINE, CAFFEINE
 ERIVEDGE, VISMODEGIB
 ERLEADA, APALUTAMIDE
 ERLOTINIB HYDROCHLORIDE, ERLOTINIB HYDROCHLORIDE
 ERRIN, NORETHINDRONE
 ERTACZO, SERTACONAZOLE NITRATE
 ERTAPENEM SODIUM, ERTAPENEM SODIUM
 ERY-TAB, ERYTHROMYCIN
 ERYC, ERYTHROMYCIN
 ERYGEL, ERYTHROMYCIN
 ERYPED, ERYTHROMYCIN ETHYLSUCCINATE
 ERYTHROCIN, ERYTHROMYCIN LACTOBIONATE
 ERYTHROCIN STEARATE, ERYTHROMYCIN STEARATE
 ERYTHROMYCIN, ERYTHROMYCIN
 ERYTHROMYCIN AND BENZOYL PEROXIDE, BENZOYL PEROXIDE
 ERYTHROMYCIN ETHYLSUCCINATE, ERYTHROMYCIN ETHYLSUCCINATE
 ESBRIET, PIRFENIDONE
 ESCITALOPRAM OXALATE, ESCITALOPRAM OXALATE
 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 AND NORGESTIMATE, ESTRADIOL
 ESTRADIOL VALERATE, ESTRADIOL VALERATE
 ESTRING, ESTRADIOL
 ESTROGEL, ESTRADIOL
 ESTROSTEP FE, ETHINYL ESTRADIOL
 ESZOPICLONE, ESZOPICLONE
 ETHACRYNATE SODIUM, ETHACRYNATE SODIUM
 ETHACRYNIC ACID, ETHACRYNIC ACID
 ETHAMBUTOL HYDROCHLORIDE, ETHAMBUTOL HYDROCHLORIDE
 ETHAMOLIN, ETHANOLAMINE OLEATE
 ETHOSUXIMIDE, ETHOSUXIMIDE
 ETHYNODIOL DIACETATE AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
 ETHYOL, AMIFOSTINE
 ETODOLAC, ETODOLAC
 ETOMIDATE, ETOMIDATE
 ETOPOPHOS PRESERVATIVE FREE, ETOPOSIDE PHOSPHATE

APPENDIX A - PRODUCT NAME INDEX

** E **

ETOPOSIDE, ETOPOSIDE
 EUCRISA, CRISABOROLE
 EURAX, CROTAMITON
 EUTHYROX, LEVOTHYROXINE SODIUM **
 EVAMIST, ESTRADIOL
 EVEKEO, AMPHETAMINE SULFATE
 EVEKEO ODT, AMPHETAMINE SULFATE
 EVEROLIMUS, EVEROLIMUS
 EVISTA, RALOXIFENE HYDROCHLORIDE
 EVOCLIN, CLINDAMYCIN PHOSPHATE
 EVOMELA, MELPHALAN HYDROCHLORIDE
 EVOTAZ, ATAZANAVIR SULFATE
 EVOXAC, CEVIMELINE HYDROCHLORIDE
 EVZIO (AUTOINJECTOR), NALOXONE HYDROCHLORIDE
 EXCEDRIN (MIGRAINE), ACETAMINOPHEN (OTC)
 EXELDERM, SULCONAZOLE NITRATE
 EXELON, RIVASTIGMINE
 EXEM FOAM KIT, AIR POLYMER-TYPE A
 EXEMESTANE, EXEMESTANE
 EXFORGE, AMLODIPINE BESYLATE
 EXFORGE HCT, AMLODIPINE BESYLATE
 EXIDINE, CHLORHEXIDINE GLUCONATE (OTC)
 EXJADE, DEFERASIROX
 EXONDYS 51, ETEPLIRSEN
 EXPAREL, BUPIVACAINE
 EXSERVAN, RILUZOLE
 EXTENDED PHENYTOIN SODIUM, PHENYTOIN SODIUM
 EXTINA, KETOCONAZOLE
 EXTRANEAL, ICODextrin
 EZALLOR, ROSUVASTATIN CALCIUM
 EZETIMIBE, EZETIMIBE
 EZETIMIBE AND ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM
 EZETIMIBE AND SIMVASTATIN, EZETIMIBE

** F **

FABIOR, TAZAROTENE
 FACTIVE, GEMIFLOXACIN MESYLATE
 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)
 FANAPT, ILOPERIDONE
 FARESTON, TOREMIFENE CITRATE
 FARXIGA, DAPAGLIFLOZIN
 FARYDAK, PANOBINOSTAT LACTATE
 FASLODEX, FULVESTRANT
 FAYOSIM, ETHINYL ESTRADIOL
 FEBUXOSTAT, FEBUXOSTAT
 FELBAMATE, FELBAMATE
 FELBATOL, FELBAMATE
 FELDENE, PIROXICAM
 FELODIPINE, FELODIPINE
 FEMARA, LETROZOLE
 FEMHRT, ETHINYL ESTRADIOL
 FEMRING, ESTRADIOL ACETATE
 FENOFIBRATE, FENOFIBRATE
 FENOFIBRATE (MICRONIZED), FENOFIBRATE
 FENOFIBRIC ACID, CHOLINE FENOFIBRATE
 FENOGLIDE, FENOFIBRATE
 FENOLDOPAM MESYLATE, FENOLDOPAM MESYLATE
 FENOPROFEN CALCIUM, FENOPROFEN CALCIUM

APPENDIX A - PRODUCT NAME INDEX

** F **

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
 FENTORA, FENTANYL CITRATE
 FERAHEME, FERUMOXYTOL
 FERRIPROX, DEFERIPRONE
 FERRLECIT, SODIUM FERRIC GLUCONATE COMPLEX
 FESOTERODINE FUMARATE, FESOTERODINE FUMARATE
 FETROJA, CEFIDEROCOL SULFATE TOSYLATE
 FETZIMA, LEVOMILNACIPRAN HYDROCHLORIDE
 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)
 FIASP, INSULIN ASPART
 FIASP FLEXTOUCH, INSULIN ASPART
 FIASP PENFILL, INSULIN ASPART
 FIBRICOR, FENOFIBRIC ACID
 FINACEA, AZELAIC ACID
 FINASTERIDE, FINASTERIDE
 FINGOLIMOD HYDROCHLORIDE, FINGOLIMOD HYDROCHLORIDE
 FIORICET W/ CODEINE, ACETAMINOPHEN
 FIORINAL, ASPIRIN
 FIORINAL W/CODEINE, ASPIRIN
 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
 FLAVORED COLESTID, COLESTIPOL HYDROCHLORIDE
 FLAVOXATE HYDROCHLORIDE, FLAVOXATE HYDROCHLORIDE
 FLECAINIDE ACETATE, FLECAINIDE ACETATE
 FLECTOR, DICLOFENAC EPOLAMINE
 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 DEXTROSE 5% IN PLASTIC CONTAINER, 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
 FLUMADINE, RIMANTADINE HYDROCHLORIDE
 FLUMAZENIL, FLUMAZENIL

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** F **

FLUNISOLIDE, FLUNISOLIDE
 FLUOCINOLONE ACETONIDE, FLUOCINOLONE ACETONIDE
 FLUOCINONIDE, FLUOCINONIDE
 FLUOCINONIDE ACETONIDE, FLUOCINOLONE ACETONIDE
 FLUOCINONIDE EMULSIFIED BASE, FLUOCINONIDE
 FLUORESCITE, FLUORESCHEIN SODIUM
 FLUORODOPA F18, FLUORODOPA F-18
 FLUOROPLEX, FLUOROURACIL
 FLUOROURACIL, FLUOROURACIL
 FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 FLUOXYMESTERONE, FLUOXYMESTERONE
 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
 FLUVASTATIN SODIUM, FLUVASTATIN SODIUM
 FLUVOXAMINE MALEATE, FLUVOXAMINE MALEATE
 FML, FLUROMETHOLONE
 FML FORTE, FLUROMETHOLONE
 FOCALIN, DEXMETHYLPHENIDATE HYDROCHLORIDE
 FOCALIN XR, DEXMETHYLPHENIDATE HYDROCHLORIDE
 FOLIC ACID, FOLIC ACID
 FOLLISTIM AQ, FOLLITROPIN ALFA/BETA
 FOLOTYN, PRALATREXATE
 FOMEPIZOLE, FOMEPIZOLE
 FONDAPARINUX SODIUM, FONDAPARINUX SODIUM
 FORANE, ISOFLURANE
 FORFIVO XL, BUPROPION HYDROCHLORIDE
 FORTAZ, CEFTAZIDIME
 FORTEO, TERIPARATIDE
 FORTESTA, TESTOSTERONE
 FOSAMAX, ALENDRONATE SODIUM
 FOSAMAX PLUS D, ALENDRONATE SODIUM
 FOSAMPRENAVIR CALCIUM, FOSAMPRENAVIR CALCIUM
 FOSAPREPITANT DIMEGLUMINE, FOSAPREPITANT DIMEGLUMINE
 FOSCAVIR, FOSCARNET SODIUM
 FOSINOPRIL SODIUM, FOSINOPRIL SODIUM
 FOSINOPRIL SODIUM AND HYDROCHLOROTHIAZIDE, FOSINOPRIL SODIUM
 FOSPHENYTOIN SODIUM, FOSPHENYTOIN SODIUM
 FOSRENOL, LANTHANUM CARBONATE
 FRAGMIN, DALTEPARIN SODIUM
 FREAMINE HBC 6.9%, AMINO ACIDS
 FREAMINE III 10%, AMINO ACIDS
 FREAMINE III 3% W/ ELECTROLYTES, AMINO ACIDS
 FREAMINE III 8.5%, AMINO ACIDS
 FREAMINE III 8.5% W/ ELECTROLYTES, AMINO ACIDS
 FROVA, FROVATRIPTAN SUCCINATE
 FROVATRIPTAN SUCCINATE, FROVATRIPTAN SUCCINATE
 FULVESTRANT, FULVESTRANT
 FURADANTIN, NITROFURANTOIN
 FUROSEMIDE, FUROSEMIDE
 FUSILEV, LEVOLEUCOVORIN CALCIUM
 FUZEON, ENFUVIRTIDE
 FYAVOLV, ETHINYL ESTRADIOL
 FYCOMPA, PERAMPANEL

** G **

GABAPENTIN, GABAPENTIN
 GABITRIL, TIAGABINE HYDROCHLORIDE
 GABLOFEN, BACLOFEN

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** G **

GADAVIST, GADOBUTROL
 GALAFOLD, MIGALASTAT HYDROCHLORIDE
 GALANTAMINE HYDROBROMIDE, GALANTAMINE HYDROBROMIDE
 GALLIUM CITRATE GA 67, GALLIUM CITRATE GA-67
 GALLIUM DOTATOC GA 68, GALLIUM DOTATOC GA-68
 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 RECOMBINANT
 GAVISCON, ALUMINUM HYDROXIDE (OTC)
 GELNIQUE, OXYBUTYNIN CHLORIDE
 GEMCITABINE HYDROCHLORIDE, GEMCITABINE HYDROCHLORIDE
 GEMFIBROZIL, GEMFIBROZIL
 GEMIFLOXACIN MESYLATE, GEMIFLOXACIN MESYLATE
 GEMZAR, GEMCITABINE HYDROCHLORIDE
 GEN-XENE, CLORAZEPATE DIPOTASSIUM
 GENERLAC, LACTULOSE
 GENGRAF, CYCLOSPORINE
 GENOPTIC, GENTAMICIN SULFATE
 GENOSYL, NITRIC OXIDE
 GENOTROPIN, SOMATROPIN
 GENOTROPIN PRESERVATIVE FREE, SOMATROPIN
 GENTAK, GENTAMICIN SULFATE
 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 1.5/30, ETHINYL ESTRADIOL
 GILDESS 1/20, ETHINYL ESTRADIOL
 GILDESS 24 FE, ETHINYL ESTRADIOL
 GILDESS FE 1.5/30, ETHINYL ESTRADIOL
 GILDESS FE 1/20, ETHINYL ESTRADIOL
 GILENYA, FINGOLIMOD HYDROCHLORIDE
 GILOTRIF, AFATINIB DIMALEATE
 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
 GLUCAGEN, GLUCAGON HYDROCHLORIDE
 GLUCAGON, GLUCAGON
 GLUCAGON, GLUCAGON HYDROCHLORIDE
 GLUCOTROL, GLIPIZIDE
 GLUCOTROL XL, GLIPIZIDE
 GLUMETZA, METFORMIN HYDROCHLORIDE
 GLYBURIDE, GLYBURIDE
 GLYBURIDE (MICRONIZED), GLYBURIDE
 GLYBURIDE AND METFORMIN HYDROCHLORIDE, GLYBURIDE
 GLYCINE 1.5% IN PLASTIC CONTAINER, GLYCINE
 GLYCOLAX, POLYETHYLENE GLYCOL 3350 (OTC)
 GLYCOPYRROLATE, GLYCOPYRROLATE

APPENDIX A - PRODUCT NAME INDEX

** G **

GLYDO, LIDOCAINE HYDROCHLORIDE
 GLYNASE, GLYBURIDE
 GLYRX-PF, GLYCOPYRROLATE
 GLYSET, MIGLITOL
 GLYXAMBI, EMPAGLIFLOZIN
 GOCOVRI, AMANTADINE HYDROCHLORIDE
 GOLYTELY, POLYETHYLENE GLYCOL 3350
 GONAL-F, FOLLITROPIN ALFA/BETA
 GONAL-F RFF, FOLLITROPIN ALFA/BETA
 GONAL-F RFF REDI-JECT, FOLLITROPIN ALFA/BETA
 GONITRO, NITROGLYCERIN
 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
 GUANIDINE HYDROCHLORIDE, GUANIDINE HYDROCHLORIDE
 GVOKE HYOPEN, GLUCAGON
 GVOKE PFS, GLUCAGON
 GYNAZOLE-1, BUTOCONAZOLE NITRATE

** H **

H.P. ACTHAR GEL, CORTICOTROPIN
 HABITROL, NICOTINE (OTC)
 HAILEY 1.5/30, 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
 HALDOL, HALOPERIDOL LACTATE
 HALOBETASOL PROPIONATE, HALOBETASOL PROPIONATE
 HALOG, HALCINONIDE
 HALOPERIDOL, HALOPERIDOL
 HALOPERIDOL, HALOPERIDOL LACTATE
 HALOPERIDOL DECANOATE, HALOPERIDOL DECANOATE
 HARVONI, LEDIPASVIR
 HEATHER, NORETHINDRONE
 HECTOROL, DOXERCALCIFEROL
 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% IN PLASTIC CONTAINER, 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
 HEPATAMINE 8%, AMINO ACIDS
 HEPSERA, ADEFOVIR DIPIVOXIL

APPENDIX A - PRODUCT NAME INDEX**** H ****

HER STYLE, LEVONORGESTREL (OTC)
HETLIOZ, 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
HUMALOG, INSULIN LISPRO RECOMBINANT
HUMALOG KWIKPEN, INSULIN LISPRO RECOMBINANT
HUMALOG MIX 50/50, INSULIN LISPRO PROTAMINE RECOMBINANT
HUMALOG MIX 50/50 KWIKPEN, INSULIN LISPRO PROTAMINE RECOMBINANT
HUMALOG MIX 75/25, INSULIN LISPRO PROTAMINE RECOMBINANT
HUMALOG MIX 75/25 KWIKPEN, INSULIN LISPRO PROTAMINE RECOMBINANT
HUMALOG TEMPO PEN, INSULIN LISPRO RECOMBINANT
HUMATROPE, SOMATROPIN
HUMULIN 70/30, INSULIN RECOMBINANT HUMAN (OTC)
HUMULIN 70/30 PEN, INSULIN RECOMBINANT HUMAN (OTC)
HUMULIN N, INSULIN SUSP ISOPHANE RECOMBINANT HUMAN (OTC)
HUMULIN R, INSULIN HUMAN
HUMULIN R, INSULIN RECOMBINANT HUMAN (OTC)
HUMULIN R KWIKPEN, INSULIN HUMAN
HUMULIN R PEN, INSULIN RECOMBINANT HUMAN (OTC)
HYCANTIN, TOPOTECAN HYDROCHLORIDE
HYCODAN, HOMATROPINE METHYLBROMIDE
HYDRA-ZIDE, HYDRALAZINE HYDROCHLORIDE
HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
HYDREA, HYDROXYUREA
HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
HYDROCODONE BITARTRATE, HYDROCODONE BITARTRATE
HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
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
HYDROCODONE POLISTIREX AND CHLORPHENIRAMNE POLISTIREX, CHLORPHENIRAMINE POLISTIREX
HYDROCORTISONE, HYDROCORTISONE
HYDROCORTISONE AND ACETIC ACID, ACETIC ACID, GLACIAL
HYDROCORTISONE BUTYRATE, HYDROCORTISONE BUTYRATE
HYDROCORTISONE IN ABSORBASE, HYDROCORTISONE
HYDROCORTISONE VALERATE, HYDROCORTISONE VALERATE
HYDROMORPHONE HYDROCHLORIDE, HYDROMORPHONE HYDROCHLORIDE
HYDROXOCOBALAMIN, HYDROXOCOBALAMIN
HYDROXYCHLOROQUINE SULFATE, HYDROXYCHLOROQUINE SULFATE
HYDROXYPROGESTERONE CAPROATE, HYDROXYPROGESTERONE CAPROATE
HYDROXYUREA, HYDROXYUREA
HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE
HYDROXYZINE PAMOATE, HYDROXYZINE PAMOATE
HYLENEX RECOMBINANT, HYALURONIDASE RECOMBINANT HUMAN
HYSINGLA ER, HYDROCODONE BITARTRATE
HYZAAR, HYDROCHLOROTHIAZIDE

**** I ****

IBANDRONATE SODIUM, IBANDRONATE SODIUM
IBRANCE, PALBOCICLIB
IBSRELA, TENAPANOR HYDROCHLORIDE
IBU-TAB, IBUPROFEN
IBU-TAB 200, IBUPROFEN (OTC)
IBUPROFEN, IBUPROFEN (OTC)
IBUPROFEN, IBUPROFEN
IBUPROFEN AND DIPHENHYDRAMINE CITRATE, DIPHENHYDRAMINE CITRATE (OTC)
IBUPROFEN AND DIPHENHYDRAMINE HYDROCHLORIDE, DIPHENHYDRAMINE HYDROCHLORIDE (OTC)
IBUPROFEN AND PHENYLEPHRINE HYDROCHLORIDE, IBUPROFEN (OTC)
IBUPROFEN AND PSEUDOEPHEDRINE HYDROCHLORIDE, IBUPROFEN (OTC)
IBUPROFEN LYSINE, IBUPROFEN LYSINE

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** I **

IBUPROFEN SODIUM, IBUPROFEN SODIUM (OTC)
 IBUPROHM COLD AND SINUS, IBUPROFEN (OTC)
 IBUTILIDE FUMARATE, IBUTILIDE FUMARATE
 IC-GREEN, INDOCYANINE GREEN
 ICATIBANT ACETATE, ICATIBANT ACETATE
 ICLEVIA, ETHINYL ESTRADIOL
 ICLUSIG, PONATINIB HYDROCHLORIDE
 IDAMYCIN PFS, IDARUBICIN HYDROCHLORIDE
 IDARUBICIN HYDROCHLORIDE, IDARUBICIN HYDROCHLORIDE
 IDARUBICIN HYDROCHLORIDE PFS, IDARUBICIN HYDROCHLORIDE
 IDHIFA, ENASIDENIB MESYLATE
 IDKIT:HP, CITRIC ACID
 IFEX, IFOSFAMIDE
 IFOSFAMIDE, IFOSFAMIDE
 ILEVRO, NEPAFENAC
 ILUVIEN, FLUOCINOLONE ACETONIDE
 IMATINIB MESYLATE, IMATINIB MESYLATE
 IMBRUVICA, IBRUTINIB
 IMIPENEM AND CILASTATIN, CILASTATIN SODIUM
 IMIPRAMINE HYDROCHLORIDE, IMIPRAMINE HYDROCHLORIDE
 IMIPRAMINE PAMOATE, IMIPRAMINE PAMOATE
 IMIQUIMOD, IMIQUIMOD
 IMITREX, SUMATRIPTAN
 IMITREX, SUMATRIPTAN SUCCINATE
 IMITREX STATDOSE, SUMATRIPTAN SUCCINATE
 IMODIUM A-D, LOPERAMIDE HYDROCHLORIDE (OTC)
 IMODIUM MULTI-SYMPATOM RELIEF, LOPERAMIDE HYDROCHLORIDE (OTC)
 IMPAVIDO, MILTEFOSINE
 IMPOYZ, CLOBETASOL PROPIONATE
 IMURAN, AZATHIOPRINE
 IMVEXXY, ESTRADIOL
 INAPSINE, DROPERIDOL
 INBRIJA, LEVODOPA
 INCASSIA, NORETHINDRONE
 INCRELEX, MECASERMIN RECOMBINANT
 INCRUSE ELLIPTA, UMECLIDINIUM BROMIDE
 INDAPAMIDE, INDAPAMIDE
 INDERAL LA, PROPRANOLOL HYDROCHLORIDE
 INDIUM IN 111 CHLORIDE, INDIUM IN-111 CHLORIDE
 INDIUM IN 111 OXYQUINOLINE, INDIUM IN-111 OXYQUINOLINE
 INDOCIN, INDOMETHACIN
 INDOCIN, INDOMETHACIN SODIUM
 INDOCYANINE GREEN, INDOCYANINE GREEN
 INDOMETHACIN, INDOMETHACIN
 INDOMETHACIN SODIUM, INDOMETHACIN SODIUM
 INFANT'S ADVIL, IBUPROFEN (OTC)
 INFANTS' FEVERALL, ACETAMINOPHEN (OTC)
 INFASURF PRESERVATIVE FREE, CALFACTANT
 INFED, IRON DEXTRAN
 INFUGEM, GEMCITABINE HYDROCHLORIDE
 INFUMORPH, MORPHINE SULFATE
 INFUVITE ADULT, ALPHA-TOCOPHEROL ACETATE
 INFUVITE PEDIATRIC, ASCORBIC ACID
 INFUVITE PEDIATRIC (PHARMACY BULK PACKAGE), ASCORBIC ACID
 INGENOL MEBUTATE, INGENOL MEBUTATE
 INGREGZA, VALBENAZINE TOSYLATE
 INJECTAFER, FERRIC CARBOXYMALTOSIDE
 INLYTA, AXITINIB
 INNOPRAN XL, PROPRANOLOL HYDROCHLORIDE
 INOMAX, NITRIC OXIDE
 INREBIC, FEDRATINIB HYDROCHLORIDE
 INSPIRA, EPLERENONE
 INTEGRILIN, EPTIFIBATIDE
 INTELENCE, ETRAVIRINE
 INTRALIPID 10%, SOYBEAN OIL

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** I **

INTRALIPID 20%, SOYBEAN OIL
 INTRALIPID 30%, SOYBEAN OIL
 INTRAROSA, PRASTERONE
 INTROVALE, ETHINYL ESTRADIOL
 INTUNIV, GUANFACINE HYDROCHLORIDE
 INVANZ, ERTAPENEM SODIUM
 INVEGA, PALIPERIDONE
 INVEGA SUSTENNA, PALIPERIDONE PALMITATE
 INVEGA TRINZA, PALIPERIDONE PALMITATE
 INVELTYS, LOTEHPREDNOL ETABONATE
 INVIRASE, SAQUINAVIR MESYLATE
 INVOKAMET, CANAGLIFLOZIN
 INVOKAMET XR, CANAGLIFLOZIN
 INVOKANA, CANAGLIFLOZIN
 IONOSOL MB AND DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
 IOPIDINE, APRACLONIDINE HYDROCHLORIDE
 IOSAT, POTASSIUM IODIDE (OTC)
 IPRATROPIUM BROMIDE, IPRATROPIUM BROMIDE
 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 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
 ISUPREL, ISOPROTERENOL HYDROCHLORIDE
 ITRACONAZOLE, ITRACONAZOLE
 IVERMECTIN, IVERMECTIN
 IVY BLOCK, BENTOQUATAM (OTC)
 IXEMPRA KIT, IXABEPILONE

** 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

APPENDIX A - PRODUCT NAME INDEX

** J **

JEANATOPE, ALBUMIN IODINATED I-125 SERUM
 JENCYCLA, NORETHINDRONE
 JENTADUETO, LINAGLIPTIN
 JENTADUETO XR, LINAGLIPTIN
 JEVTANA KIT, CABAZITAXEL
 JORNAY PM, METHYLPHENIDATE HYDROCHLORIDE
 JUBLIA, EFINAACONAZOLE
 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 IBUPROFEN, IBUPROFEN (OTC)
 JUNIOR STRENGTH MOTRIN, IBUPROFEN (OTC)
 JUXTAPID, LOMITAPIDE MESYLATE
 JYNARQUE, TOLVAPTAN

** K **

K-TAB, POTASSIUM CHLORIDE
 KABIVEN IN PLASTIC CONTAINER, AMINO ACIDS
 KADIAN, MORPHINE SULFATE
 KAITLIB FE, ETHINYL ESTRADIOL
 KALETRA, LOPINAVIR
 KALEXATE, SODIUM POLYSTYRENE SULFONATE
 KALLIGA, DESOGESTREL
 KALYDECO, IVACAFTOR
 KAPSPARGO SPRINKLE, METOPROLOL SUCCINATE
 KAPVAY, CLONIDINE HYDROCHLORIDE
 KARBINAL ER, CARBINOXAMINE MALEATE
 KARIVA, DESOGESTREL
 KATERZIA, AMLODIPINE BENZOATE
 KAZANO, ALOGLIPTIN BENZOATE
 KEFLEX, CEPHALEXIN
 KELNOR, ETHINYL ESTRADIOL
 KEMEYA, DROSPIRENONE
 KENALOG, TRIAMCINOLONE ACETONIDE
 KENALOG-10, TRIAMCINOLONE ACETONIDE
 KENALOG-40, TRIAMCINOLONE ACETONIDE
 KENALOG-80, TRIAMCINOLONE ACETONIDE
 KENGREAL, CANGRELOR
 KEPPRA, LEVETIRACETAM
 KEPPRA XR, LEVETIRACETAM
 KERYDIN, TAVABOROLE
 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)
 KETOZOLE, KETOCONAZOLE
 KEVEYIS, DICHLORPHENAMIDE
 KHAPZORY, LEVOLEUCOVORIN
 KIMIDESS, DESOGESTREL
 KINEVAC, SINCALIDE
 KIONEX, SODIUM POLYSTYRENE SULFONATE
 KISQALI, RIBOCICLIB SUCCINATE
 KISQALI FEMARA CO-PACK (COPACKAGED), LETROZOLE
 KITABIS PAK, TOBRAMYCIN
 KLARON, SULFACETAMIDE SODIUM
 KLONOPIN, CLONAZEPAM
 KLOR-CON, POTASSIUM CHLORIDE
 KLOR-CON M10, POTASSIUM CHLORIDE
 KLOR-CON M15, POTASSIUM CHLORIDE

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** K **

KLOR-CON M20, POTASSIUM CHLORIDE
 KOMBIGLYZE XR, METFORMIN HYDROCHLORIDE
 KORLYM, MIFEPRISTONE
 KOVANAZE, OXYMETAZOLINE HYDROCHLORIDE
 KRINTAFEL, TAFENOQUINE SUCCINATE
 KURVELO, ETHINYL ESTRADIOL
 KUVAN, SAPROPTERIN DIHYDROCHLORIDE
 KYBELLA, DEOXYCHOLIC ACID
 KYLEENA, LEVONORGESTREL
 KYPROLIS, CARFILZOMIB
 KYRA, DROSPIRENONE

** L **

LABETALOL HYDROCHLORIDE, LABETALOL HYDROCHLORIDE
 LACRISERT, HYDROXYPROPYL CELLULOSE
 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, TERBINAFFINE HYDROCHLORIDE (OTC)
 LAMISIL, TERBINAFFINE HYDROCHLORIDE
 LAMISIL AT, TERBINAFFINE (OTC)
 LAMISIL AT, TERBINAFFINE HYDROCHLORIDE (OTC)
 LAMIVUDINE, LAMIVUDINE
 LAMIVUDINE AND ZIDOVUDINE, LAMIVUDINE
 LAMOTRIGINE, LAMOTRIGINE
 LANORINAL, ASPIRIN
 LANOXIN, DIGOXIN
 LANOXIN PEDIATRIC, DIGOXIN
 LANSOPRAZOLE, LANSOPRAZOLE (OTC)
 LANSOPRAZOLE, LANSOPRAZOLE
 LANSOPRAZOLE, AMOXICILLIN AND CLARITHROMYCIN, AMOXICILLIN
 LANTHANUM CARBONATE, LANTHANUM CARBONATE
 LANTUS, INSULIN GLARGINE RECOMBINANT
 LANTUS SOLOSTAR, INSULIN GLARGINE RECOMBINANT
 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
 LAROTID, AMOXICILLIN
 LARYNG-O-JET KIT, LIDOCAINE HYDROCHLORIDE
 LASIX, FUROSEMIDE
 LASTACAFT, ALCAFTADINE
 LATANOPROST, LATANOPROST
 LATISSE, BIMATOPROST
 LATUDA, LURASIDONE HYDROCHLORIDE
 LAX-LYTE WITH FLAVOR PACKS, POLYETHYLENE GLYCOL 3350
 LAZANDA, FENTANYL CITRATE
 LEFLUNOMIDE, LEFLUNOMIDE
 LENVIMA, LENVATINIB MESYLATE
 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
 LEVALBUTEROL HYDROCHLORIDE, LEVALBUTEROL HYDROCHLORIDE
 LEVEMIR, INSULIN DETEMIR RECOMBINANT

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** L **

LEVEMIR FLEXTOUCH, INSULIN DETEMIR RECOMBINANT
 LEVETIRACETAM, LEVETIRACETAM
 LEVETIRACETAM IN SODIUM CHLORIDE, LEVETIRACETAM
 LEVITRA, VARDENAFIL HYDROCHLORIDE
 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
 LEVOLEUCOVORIN CALCIUM, LEVOLEUCOVORIN CALCIUM
 LEVONEST, ETHINYL ESTRADIOL
 LEVONORGESTREL, LEVONORGESTREL (OTC)
 LEVONORGESTREL, LEVONORGESTREL
 LEVONORGESTREL AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
 LEVONORGESTREL AND ETHINYL ESTRADIOL AND ETHINYL ESTRADIOL, 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
 LEXIVA, FOSAMPRENAVIR CALCIUM
 LIALDA, MESALAMINE
 LIBRAX, CHLORDIAZEPOXIDE HYDROCHLORIDE
 LIBRIUM, CHLORDIAZEPOXIDE HYDROCHLORIDE
 LIDEX, FLUOCINONIDE
 LIDEX-E, FLUOCINONIDE
 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
 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
 LILETTA, LEVONORGESTREL
 LINCOCIN, LINCOMYCIN HYDROCHLORIDE
 LINCOMYCIN, LINCOMYCIN HYDROCHLORIDE
 LINDANE, LINDANE
 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
 LISINOPRIL, LISINOPRIL

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** L **

LISINOPRIL AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 LITHIUM CARBONATE, LITHIUM CARBONATE
 LITHIUM CITRATE, LITHIUM CITRATE
 LITHOBID, LITHIUM CARBONATE
 LITHOSTAT, ACETOHYDROXAMIC ACID
 LIVALO, PITAVASTATIN CALCIUM
 LO LOESTRIN FE, ETHINYL ESTRADIOL
 LO SIMPESSE, ETHINYL ESTRADIOL
 LO-MALMOREDE, ETHINYL ESTRADIOL
 LO-ZUMANDIMINE, DROSPIRENONE
 LOCROID, HYDROCORTISONE BUTYRATE
 LOCROID LIPOCREAM, HYDROCORTISONE BUTYRATE
 LODOSYN, CARBIDOPA
 LOESTRIN 21 1.5/30, ETHINYL ESTRADIOL
 LOESTRIN 21 1/20, ETHINYL ESTRADIOL
 LOESTRIN 24 FE, ETHINYL ESTRADIOL
 LOESTRIN FE 1.5/30, ETHINYL ESTRADIOL
 LOESTRIN FE 1/20, ETHINYL ESTRADIOL
 LOGILIA, ULIPRISTAL ACETATE
 LOKELMA, SODIUM ZIRCONIUM CYCLOSILICATE
 LOMAIRA, PHENTERMINE HYDROCHLORIDE
 LOMOTIL, ATROPINE SULFATE
 LONHALA MAGNAIR KIT, GLYCOPYRROLATE
 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)
 LORATADINE REDIDOSE, LORATADINE (OTC)
 LORAZEPAM, LORAZEPAM
 LORAZEPAM INTENSOL, LORAZEPAM
 LORBRENA, LORLATINIB
 LORYNA, DROSPIRENONE
 LOSARTAN POTASSIUM, LOSARTAN POTASSIUM
 LOSARTAN POTASSIUM AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 LOSEASONIQUE, ETHINYL ESTRADIOL
 LOTEMAX, LOTEPIREDNOL ETABONATE
 LOTEMAX SM, LOTEPIREDNOL ETABONATE
 LOTENSIN, BENAZEPRIL HYDROCHLORIDE
 LOTENSIN HCT, BENAZEPRIL HYDROCHLORIDE
 LOTEPIREDNOL ETABONATE, LOTEPIREDNOL ETABONATE
 LOTREL, AMLODIPINE BESYLATE
 LOTRIMIN ULTRA, BUTENAFINE HYDROCHLORIDE (OTC)
 LOTRISONE, BETAMETHASONE DIPROPIONATE
 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
 LUCEMYRA, LOFEXIDINE HYDROCHLORIDE
 LUMASON, SULFUR HEXAFLUORIDE LIPID-TYPE A MICROSPHERES
 LUMIFY, BRIMONIDINE TARTRATE (OTC)
 LUMIGAN, BIMATOPROST
 LUNESTA, ESZOPICLONE
 LUPANETA PACK, LEUPROLIDE ACETATE
 LUPRON DEPOT, LEUPROLIDE ACETATE

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LUPRON DEPOT-PED, LEUPROLIDE ACETATE
 LURASIDONE HYDROCHLORIDE, LURASIDONE HYDROCHLORIDE
 LUTATHERA, LUTETIUM DOTATATE LU-177
 LUVOX, FLUVOXAMINE MALEATE
 LUXIQ, BETAMETHASONE VALERATE
 LUZU, LULICONAZOLE
 LYMPHOSEEK KIT, TECHNETIUM TC-99M TILMANOCEPT
 LYNPARZA, OLAPARIB
 LYRICA, PREGABALIN
 LYRICA CR, PREGABALIN
 LYSODREN, MITOTANE
 LYSTEDA, TRANEXAMIC ACID

**** M ****

M-ZOLE 3 COMBINATION PACK, MICONAZOLE NITRATE (OTC)
 M.V.I. ADULT, ASCORBIC ACID
 M.V.I. ADULT (PHARMACY BULK PACKAGE), ASCORBIC ACID
 M.V.I. PEDIATRIC, ASCORBIC ACID
 MACRILEN, MACIMORELIN ACETATE
 MACROBID, NITROFURANTOIN
 MACRODANTIN, NITROFURANTOIN, MACROCRYSTALLINE
 MACUGEN, PEGAPTANIB SODIUM
 MAFENIDE ACETATE, MAFENIDE ACETATE
 MAGNESIUM SULFATE, MAGNESIUM SULFATE
 MAGNESIUM SULFATE IN DEXTROSE 5% IN PLASTIC CONTAINER, MAGNESIUM SULFATE
 MAGNESIUM SULFATE IN PLASTIC CONTAINER, MAGNESIUM SULFATE
 MAKENA, HYDROXYPROGESTERONE CAPROATE
 MAKENA (AUTOINJECTOR), HYDROXYPROGESTERONE CAPROATE
 MAKENA PRESERVATIVE FREE, HYDROXYPROGESTERONE CAPROATE
 MALARONE, ATOVAQUONE
 MALARONE PEDIATRIC, ATOVAQUONE
 MALATHION, MALATHION
 MALMOREDE, ETHINYL ESTRADIOL
 MANGANESE CHLORIDE IN PLASTIC CONTAINER, MANGANESE CHLORIDE
 MANNITOL 10% IN PLASTIC CONTAINER, MANNITOL
 MANNITOL 15% IN PLASTIC CONTAINER, MANNITOL
 MANNITOL 20% IN PLASTIC CONTAINER, MANNITOL
 MANNITOL 25%, MANNITOL
 MANNITOL 5% IN PLASTIC CONTAINER, MANNITOL
 MAPROTILINE HYDROCHLORIDE, MAPROTILINE HYDROCHLORIDE
 MARCAINE, BUPIVACAINE HYDROCHLORIDE
 MARCAINE HYDROCHLORIDE, BUPIVACAINE HYDROCHLORIDE
 MARCAINE HYDROCHLORIDE PRESERVATIVE FREE, BUPIVACAINE HYDROCHLORIDE
 MARCAINE HYDROCHLORIDE W/ EPINEPHRINE, BUPIVACAINE HYDROCHLORIDE
 MARCAINE HYDROCHLORIDE W/ EPINEPHRINE PRESERVATIVE FREE, BUPIVACAINE HYDROCHLORIDE
 MARINOL, DRONABINOL
 MARLISSA, ETHINYL ESTRADIOL
 MARPLAN, ISOCARBOXAZID
 MARQIBO KIT, VINCRISTINE SULFATE
 MATULANE, PROCARBAZINE HYDROCHLORIDE
 MAVENCLAD, CLADRIBINE
 MAVYRET, GLECAPREVIR
 MAXALT, RIZATRIPTAN BENZOATE
 MAXALT-MLT, RIZATRIPTAN BENZOATE
 MAXIDEX, DEXAMETHASONE
 MAXIPIPE, CEFEPIME HYDROCHLORIDE
 MAXITROL, DEXAMETHASONE
 MAXZIDE, HYDROCHLOROTHIAZIDE
 MAXZIDE-25, HYDROCHLOROTHIAZIDE
 MAYZENT, SIPONIMOD FUMARIC ACID
 MD-GASTROVIEW, DIATRIZOATE MEGLUMINE
 MECAMYLAMINE HYDROCHLORIDE, MECAMYLAMINE HYDROCHLORIDE
 MECLIZINE HYDROCHLORIDE, MECLIZINE HYDROCHLORIDE
 MECLOFENAMATE SODIUM, MECLOFENAMATE SODIUM
 MEDROL, METHYLPREDNISOLONE

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** M **

MEDROXYPROGESTERONE ACETATE, MEDROXYPROGESTERONE ACETATE
 MEFENAMIC ACID, MEFENAMIC ACID
 MEFLOQUINE HYDROCHLORIDE, MEFLOQUINE HYDROCHLORIDE
 MEFOXIN IN PLASTIC CONTAINER, CEFOXITIN SODIUM
 MEGACE ES, MEGESTROL ACETATE
 MEGATOPE, ALBUMIN IODINATED I-131 SERUM
 MEGESTROL ACETATE, MEGESTROL ACETATE
 MEKINIST, TRAMETINIB DIMETHYL SULFOXIDE
 MEKTOVI, BINIMETINIB
 MELAMISA, DROSPIRENONE
 MELOXICAM, MELOXICAM
 MELPHALAN, MELPHALAN
 MELPHALAN HYDROCHLORIDE, MELPHALAN HYDROCHLORIDE
 MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
 MEMBRANEBLUE, TRYPAN BLUE
 MEN'S ROGAINE, MINOXIDIL (OTC)
 MENEST, ESTROGENS, ESTERIFIED
 MENOPUR, MENOTROPINS (FSH)
 MENOSTAR, ESTRADIOL
 MENTAX, BUTENAFINE HYDROCHLORIDE
 MEPERIDINE HYDROCHLORIDE, MEPERIDINE HYDROCHLORIDE
 MEPERIDINE HYDROCHLORIDE PRESERVATIVE FREE, MEPERIDINE HYDROCHLORIDE
 MEPHYTON, PHYTONADIONE
 MEPROBAMATE, MEPROBAMATE
 MEPRON, ATOVAQUONE
 MERCAPTOPYRINE, MERCAPTOPYRINE
 MEROPENEM, MEROPENEM
 MEROPENEM AND SODIUM CHLORIDE IN DUPLEX CONTAINER, MEROPENEM
 MERREM, MEROPENEM
 MESALAMINE, MESALAMINE
 MESNA, MESNA
 MESNEX, MESNA
 MESTINON, PYRIDOSTIGMINE BROMIDE
 METADATE CD, METHYLPHENIDATE HYDROCHLORIDE
 METAPROTERENOL SULFATE, METAPROTERENOL SULFATE
 METASTRON, STRONTIUM CHLORIDE SR-89
 METAXALONE, METAXALONE
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 METHADONE HYDROCHLORIDE, METHADONE HYDROCHLORIDE
 METHADONE HYDROCHLORIDE INTENSOL, METHADONE HYDROCHLORIDE
 METHADOSE, METHADONE HYDROCHLORIDE
 METHAMPHETAMINE HYDROCHLORIDE, METHAMPHETAMINE HYDROCHLORIDE
 METHAZOLAMIDE, METHAZOLAMIDE
 METHENAMINE HIPPURATE, METHENAMINE HIPPURATE
 METHERGINE, METHYLERGONOVINE MALEATE
 METHIMAZOLE, METHIMAZOLE
 METHOCARBAMOL, METHOCARBAMOL
 METHOCARBAMOL AND ASPIRIN, ASPIRIN
 METHOTREXATE PRESERVATIVE FREE, METHOTREXATE SODIUM
 METHOTREXATE SODIUM, METHOTREXATE SODIUM
 METHOTREXATE SODIUM PRESERVATIVE FREE, METHOTREXATE SODIUM
 METHOXSALEN, METHOXSALEN
 METHSCOPOLAMINE BROMIDE, METHSCOPOLAMINE BROMIDE
 METHYLDOPA, METHYLDOPA
 METHYLDOPA AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 METHYLERGONOVINE MALEATE, METHYLERGONOVINE MALEATE
 METHYLIN, METHYLPHENIDATE HYDROCHLORIDE
 METHYLIN ER, METHYLPHENIDATE HYDROCHLORIDE
 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

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** M **

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
 MEXILETINE HYDROCHLORIDE, MEXILETINE HYDROCHLORIDE
 MIACALCIN, CALCITONIN SALMON
 MIBELAS 24 FE, ETHINYL ESTRADIOL
 MICAFUNGIN SODIUM, MICAFUNGIN 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
 MICRO-K, POTASSIUM CHLORIDE
 MICRO-K 10, POTASSIUM CHLORIDE
 MICROGESTIN 1.5/30, ETHINYL ESTRADIOL
 MICROGESTIN 1/20, ETHINYL ESTRADIOL
 MICROGESTIN FE 1.5/30, ETHINYL ESTRADIOL
 MICROGESTIN FE 1/20, ETHINYL ESTRADIOL
 MICROZIDE, HYDROCHLOROTHIAZIDE
 MIDAMOR, AMILORIDE HYDROCHLORIDE
 MIDAZOLAM HYDROCHLORIDE, MIDAZOLAM HYDROCHLORIDE
 MIDAZOLAM HYDROCHLORIDE PRESERVATIVE FREE, MIDAZOLAM HYDROCHLORIDE
 MIDODRINE HYDROCHLORIDE, MIDODRINE HYDROCHLORIDE
 MIDOL LIQUID GELS, IBUPROFEN (OTC)
 MIFEPREX, MIFEPRISTONE
 MIFEPRISTONE, MIFEPRISTONE
 MIGERGOT, CAFFEINE
 MIGLITOL, MIGLITOL
 MIGLUSTAT, MIGLUSTAT
 MIGRANAL, DIHYDROERGOTAMINE MESYLATE
 MILI, ETHINYL ESTRADIOL
 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
 MINASTRIN 24 FE, ETHINYL ESTRADIOL
 MINIPRESS, PRAZOSIN HYDROCHLORIDE
 MINIRIN, DESMOPRESSIN ACETATE
 MINIVELLE, ESTRADIOL
 MINOCIN, MINOCYCLINE HYDROCHLORIDE
 MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE
 MINOLIRA, MINOCYCLINE HYDROCHLORIDE
 MINOXIDIL, MINOXIDIL (OTC)
 MINOXIDIL, MINOXIDIL
 MINOXIDIL (FOR MEN), MINOXIDIL (OTC)
 MINOXIDIL (FOR WOMEN), MINOXIDIL (OTC)
 MINOXIDIL EXTRA STRENGTH (FOR MEN), MINOXIDIL (OTC)
 MIOCHOL-E, ACETYLCHOLINE CHLORIDE
 MIOSTAT, CARBACHOL
 MIRALAX, POLYETHYLENE GLYCOL 3350 (OTC)
 MIRAPEX, PRAMIPEXOLE DIHYDROCHLORIDE
 MIRAPEX ER, PRAMIPEXOLE DIHYDROCHLORIDE

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** M **

MIRENA, LEVONORGESTREL
MIRTAZAPINE, MIRTAZAPINE
MIRVASO, BRIMONIDINE TARTRATE
MISOPROSTOL, MISOPROSTOL
MITIGARE, COLCHICINE
MITIGO, MORPHINE SULFATE
MITOMYCIN, MITOMYCIN
MITOSOL, MITOMYCIN
MITOXANTRONE HYDROCHLORIDE, MITOXANTRONE HYDROCHLORIDE
MOBIC, MELOXICAM
MODAFINIL, MODAFINIL
MOEXIPRIL HYDROCHLORIDE, MOEXIPRIL HYDROCHLORIDE
MOEXIPRIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
MOLINDONE HYDROCHLORIDE, MOLINDONE HYDROCHLORIDE
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
MONOFERRIC, FERRIC DERISOMALTOSE
MONOKET, ISOSORBIDE MONONITRATE
MONTELUKAST SODIUM, MONTELUKAST SODIUM
MONUROL, FOSFOMYCIN TROMETHAMINE
MORPHABOND ER, MORPHINE SULFATE
MORPHINE SULFATE, MORPHINE SULFATE
MOTEGRITY, PRUCALOPRIDE SUCCINATE
MOTOFEN, ATROPINE SULFATE
MOTRIN IB, IBUPROFEN (OTC)
MOVANTIK, NALOXEGOL OXALATE
MOVIPREP, ASCORBIC ACID
MOXEZA, MOXIFLOXACIN HYDROCHLORIDE
MOXIDECTIN, MOXIDECTIN
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
MUPIROCIN, MUPIROCIN
MUPIROCIN, MUPIROCIN CALCIUM
MUSE, ALPROSTADIL
MYAMBUTOL, ETHAMBUTOL HYDROCHLORIDE
MYCAMINE, MICAFUNGIN SODIUM
MYCOBUTIN, RIFABUTIN
MYCOPHENOLATE MOFETIL, MYCOPHENOLATE MOFETIL
MYCOPHENOLATE MOFETIL HYDROCHLORIDE, MYCOPHENOLATE MOFETIL HYDROCHLORIDE
MYCOPHENOLIC ACID, MYCOPHENOLIC ACID
MYDAYIS, AMPHETAMINE ASPARTATE
MYDRIACYL, TROPICAMIDE
MYFORTIC, MYCOPHENOLIC ACID
MYKACET, NYSTATIN
MYLERAN, BUSULFAN
MYORISAN, ISOTRETINOIN
MYOVIEW 30ML, TECHNETIUM TC-99M TETROFOSMIN KIT
MYRBETRIQ, MIRABEGRON

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** M **

MYSOLINE, PRIMIDONE
 MYTESI, CROFELEMER
 MYXREDLIN, INSULIN HUMAN
 MYZILRA, ETHINYL ESTRADIOL

** N **

NABUMETONE, NABUMETONE
 NADOLOL, NADOLOL
 NADOLOL AND BENDROFLUMETHIAZIDE, BENDROFLUMETHIAZIDE
 NAFACILLIN SODIUM, NAFACILLIN SODIUM
 NAFTIFINE HYDROCHLORIDE, NAFTIFINE HYDROCHLORIDE
 NAFTIN, NAFTIFINE HYDROCHLORIDE
 NALBUPHINE HYDROCHLORIDE, NALBUPHINE HYDROCHLORIDE
 NALFON, FENOPROFEN CALCIUM
 NALLPEN IN PLASTIC CONTAINER, NAFACILLIN SODIUM
 NALOXONE, NALOXONE HYDROCHLORIDE
 NALOXONE HYDROCHLORIDE, NALOXONE HYDROCHLORIDE
 NALOXONE HYDROCHLORIDE AND PENTAZOCINE HYDROCHLORIDE, NALOXONE HYDROCHLORIDE
 NALTREXONE HYDROCHLORIDE, NALTREXONE HYDROCHLORIDE
 NAMENDA, MEMANTINE HYDROCHLORIDE
 NAMENDA XR, 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)
 NARATRIPTAN, NARATRIPTAN HYDROCHLORIDE
 NARCAN, NALOXONE HYDROCHLORIDE
 NARDIL, PHENELZINE SULFATE
 NAROPIN, ROPIVACAINE HYDROCHLORIDE
 NASACORT ALLERGY 24 HOUR, TRIAMCINOLONE ACETONIDE (OTC)
 NASCOBAL, CYANOCOBALAMIN
 NASONEX, MOMETASONE FUROATE
 NATACYN, NATAMYCIN
 NATAZIA, DIENOGEST
 NATEGLINIDE, NATEGLINIDE
 NATESTO, TESTOSTERONE
 NATROBA, SPINOSAD
 NAYZILAM, MIDAZOLAM
 NEBUPENT, PENTAMIDINE ISETHIONATE
 NEDOCROMIL SODIUM, NEDOCROMIL SODIUM
 NEFAZODONE HYDROCHLORIDE, NEFAZODONE HYDROCHLORIDE
 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
 NEOPAP, ACETAMINOPHEN (OTC)
 NEOPROFEN, IBUPROFEN LYSINE
 NEORAL, CYCLOSPORINE
 NEOSPORIN, BACITRACIN ZINC
 NEOSPORIN, GRAMICIDIN
 NEOSPORIN G.U. IRRIGANT, NEOMYCIN SULFATE
 NEOSTIGMINE METHYLSULFATE, NEOSTIGMINE METHYLSULFATE
 NEPHRAMINE 5.4%, AMINO ACIDS

APPENDIX A - PRODUCT NAME INDEX

** N **

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
 NEXIUM, ESOMEPRAZOLE MAGNESIUM
 NEXIUM 24HR, ESOMEPRAZOLE MAGNESIUM (OTC)
 NEXIUM IV, ESOMEPRAZOLE SODIUM
 NEXLETOL, BEMPEDOIC ACID
 NEXLIZET, BEMPEDOIC ACID
 NEXPLANON, ETNOGESTREL
 NEXTERONE, AMIODARONE HYDROCHLORIDE
 NIACIN, NIACIN
 NIACOR, NIACIN
 NIASPAN, NIACIN
 NICARDIPINE HYDROCHLORIDE, NICARDIPINE HYDROCHLORIDE
 NICODERM CQ, NICOTINE (OTC)
 NICORETTE, NICOTINE POLACRILEX (OTC)
 NICORETTE (MINT), NICOTINE POLACRILEX (OTC)
 NICOTINE, NICOTINE (OTC)
 NICOTINE POLACRILEX, NICOTINE POLACRILEX (OTC)
 NICOTROL, NICOTINE
 NIFEDIPINE, NIFEDIPINE
 NIKKI, DROSPIRENONE
 NILANDRON, NILUTAMIDE
 NILUTAMIDE, NILUTAMIDE
 NIMBEX, CISATRACURIUM BESYLATE
 NIMBEX PRESERVATIVE FREE, CISATRACURIUM BESYLATE
 NIMODIPINE, NIMODIPINE
 NINLARO, IXAZOMIB CITRATE
 NIPENT, PENTOSTATIN
 NIPRIDE RTU IN SODIUM CHLORIDE 0.9%, SODIUM NITROPRUSSIDE
 NISOLDIPINE, NISOLDIPINE
 NITHIODOTE, 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
 NITROPRESS, SODIUM NITROPRUSSIDE
 NITROSTAT, NITROGLYCERIN
 NITYR, NITISINONE
 NIX, PERMETHRIN (OTC)
 NIZATIDINE, NIZATIDINE
 NIZORAL, KETOCONAZOLE
 NIZORAL A-D, KETOCONAZOLE (OTC)
 NOCDURNA, DESMOPRESSIN ACETATE
 NOR-QD, NORETHINDRONE
 NORCO, ACETAMINOPHEN
 NORDITROPIN FLEXPRO, SOMATROPIN
 NOREPINEPHRINE BITARTRATE, NOREPINEPHRINE BITARTRATE
 NORETHINDRONE, NORETHINDRONE
 NORETHINDRONE ACETATE, NORETHINDRONE ACETATE

APPENDIX A - PRODUCT NAME INDEX

** N **

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
NORINYL 1+35 21-DAY, ETHINYL ESTRADIOL
NORINYL 1+35 28-DAY, ETHINYL ESTRADIOL
NORITATE, METRONIDAZOLE
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
NOURESS, CYSTEINE HYDROCHLORIDE
NOURIANZ, ISTRADÉFYLLINE
NOVOLIN 70/30, INSULIN RECOMBINANT HUMAN (OTC)
NOVOLIN N, INSULIN SUSP ISOPHANE RECOMBINANT HUMAN (OTC)
NOVOLIN R, INSULIN RECOMBINANT HUMAN (OTC)
NOVOLOG, INSULIN ASPART RECOMBINANT
NOVOLOG FLEXPEN, INSULIN ASPART RECOMBINANT
NOVOLOG MIX 70/30, INSULIN ASPART PROTAMINE RECOMBINANT
NOVOLOG MIX 70/30 FLEXPEN, INSULIN ASPART PROTAMINE RECOMBINANT
NOVOLOG PENFILL, INSULIN ASPART RECOMBINANT
NOXAFIL, POSACONAZOLE
NOXIVENT, NITRIC OXIDE
NUBEQA, DAROLUTAMIDE
NUCYNTA, TAPENTADOL HYDROCHLORIDE
NUCYNTA ER, TAPENTADOL HYDROCHLORIDE
NUEDEXTA, DEXTROMETHORPHAN HYDROBROMIDE
NULYTELY, POLYETHYLENE GLYCOL 3350
NULYTELY-FLAVORED, POLYETHYLENE GLYCOL 3350
NUMBRINO, COCAINE HYDROCHLORIDE
NUPLAZID, PIMAVANSERIN TARTRATE
NURTEC ODT, RIMEGEPANT SULFATE
NUTRILIPID 10%, SOYBEAN OIL
NUTRILIPID 20%, SOYBEAN OIL
NUTROPIN AQ NUSPIN, SOMATROPIN
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, OBETICHOIC ACID
OCTREOSCAN, INDIUM IN-111 PENTETREOTIDE KIT
OCTREOTIDE ACETATE, OCTREOTIDE ACETATE
OCTREOTIDE ACETATE (PRESERVATIVE FREE), OCTREOTIDE ACETATE

APPENDIX A - PRODUCT NAME INDEX

** O **

OCUFEN, FLURBIPROFEN SODIUM
 OCUFLOX, OFLOXACIN
 ODEFSEY, EMTRICITABINE
 ODOMZO, SONIDEGIB PHOSPHATE
 OFEV, NINTEDANIB ESYLATE
 OFIRMEV, ACETAMINOPHEN
 OFLOXACIN, OFLOXACIN
 OGEN 5, ESTROPIPATE
 OGESTREL 0.5/50-28, ETHINYL ESTRADIOL
 OLANZAPINE, OLANZAPINE
 OLANZAPINE AND FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 OLMESARTAN MEDOXOMIL, OLMESARTAN MEDOXOMIL
 OLMESARTAN MEDOXOMIL AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 OLMESARTAN MEDOXOMIL, AMLODIPINE AND HYDROCHLOROTHIAZIDE, AMLODIPINE BESYLATE
 OLOPATADINE HYDROCHLORIDE, OLOPATADINE HYDROCHLORIDE
 OLUMIANT, BARICITINIB
 OLUX, CLOBETASOL PROPIONATE
 OLUX E, CLOBETASOL PROPIONATE
 OMEGA-3-ACID ETHYL ESTERS, OMEGA-3-ACID ETHYL ESTERS
 OMEGAVEN, FISH OIL TRIGLYCERIDES
 OMEPRAZOLE, OMEPRAZOLE (OTC)
 OMEPRAZOLE, OMEPRAZOLE
 OMEPRAZOLE AND CLARITHROMYCIN AND AMOXICILLIN, AMOXICILLIN
 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
 OMNITROPE, SOMATROPIN
 ONDANSETRON, ONDANSETRON
 ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
 ONDANSETRON HYDROCHLORIDE PRESERVATIVE FREE, ONDANSETRON HYDROCHLORIDE
 ONEXTON, BENZOYL PEROXIDE
 ONFI, CLOBAZAM
 ONGLYZA, SAXAGLIPTIN HYDROCHLORIDE
 ONIVYDE, IRINOTECAN HYDROCHLORIDE
 ONMEL, ITRACONAZOLE
 ONPATTRO, PATISIRAN SODIUM
 ONZETRA XSAIL, SUMATRIPTAN SUCCINATE
 OPCICON ONE-STEP, LEVONORGESTREL (OTC)
 OPCON-A, NAPHAZOLINE HYDROCHLORIDE (OTC)
 OPSUMIT, MACITENTAN
 OPTIRAY 240, IOVERSOL
 OPTIRAY 300, IOVERSOL
 OPTIRAY 320, IOVERSOL
 OPTIRAY 350, IOVERSOL
 OPTISON, ALBUMIN HUMAN
 ORABLOC, ARTICAINE HYDROCHLORIDE
 ORACEA, DOXYCYCLINE
 ORALTAG, IOHEXOL
 ORAPRED ODT, PREDNISOLONE SODIUM PHOSPHATE
 ORAQIX, LIDOCAINE
 ORAVERSE, PHENTOLAMINE MESYLATE
 ORAVIG, MICONAZOLE
 ORBACTIV, ORITAVANCIN DIPHOSPHATE
 ORENITRAM, TREPROSTINIL DIOLAMINE

APPENDIX A - PRODUCT NAME INDEX

** O **

ORFADIN, NITISINONE
 ORILISSA, ELAGOLIX SODIUM
 ORKAMBI, IVACAFTOR
 ORPHENADRINE CITRATE, ORPHENADRINE CITRATE
 ORSYTHIA, ETHINYL ESTRADIOL
 ORTHO TRI-CYCLEN, ETHINYL ESTRADIOL
 ORTHO TRI-CYCLEN LO, ETHINYL ESTRADIOL
 ORTHO-NOVUM 7/7/7-28, ETHINYL ESTRADIOL
 ORTIKOS, BUDESONIDE
 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
 OSMOLEX ER, AMANTADINE HYDROCHLORIDE
 OSMOPREP, SODIUM PHOSPHATE, DIBASIC, ANHYDROUS
 OSPHENA, OSPEMIFENE
 OTEZLA, APREMILAST
 OTICAIR, HYDROCORTISONE
 OTIPRIO, CIPROFLOXACIN
 OTOVEL, CIPROFLOXACIN HYDROCHLORIDE
 OTREXUP, METHOTREXATE
 OVIDREL, CHORIOGONADOTROPIN ALFA
 OXACILLIN SODIUM, OXACILLIN SODIUM
 OXALIPLATIN, OXALIPLATIN
 OXANDROLONE, OXANDROLONE
 OXAPROZIN, OXAPROZIN
 OXAYDO, OXYCODONE HYDROCHLORIDE
 OXAZEPAM, OXAZEPAM
 OXBRYTA, VOXELOTOR
 OXCARBAZEPINE, OXCARBAZEPINE
 OXICONAZOLE NITRATE, OXICONAZOLE NITRATE
 OXISTAT, OXICONAZOLE NITRATE
 OXSORALEN-ULTRA, METHOXSALEN
 OXTELLAR XR, OXCARBAZEPINE
 OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE
 OXYCET, ACETAMINOPHEN
 OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN
 OXYCODONE AND ASPIRIN, ASPIRIN
 OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
 OXYCODONE HYDROCHLORIDE AND IBUPROFEN, IBUPROFEN
 OXYCONTIN, OXYCODONE HYDROCHLORIDE
 OXYMORPHONE HYDROCHLORIDE, OXYMORPHONE HYDROCHLORIDE
 OXYTOCIN, OXYTOCIN
 OXYTROL, OXYBUTYNIN
 OXYTROL FOR WOMEN, OXYBUTYNIN (OTC)
 OZEMPIC, SEMAGLUTIDE
 OZOBAX, BACLOFEN
 OZURDEX, DEXAMETHASONE

** P **

PACERONE, AMIODARONE HYDROCHLORIDE
 PACITAXEL, PACLITAXEL
 PACLITAXEL, PACLITAXEL
 PALIPERIDONE, PALIPERIDONE
 PALONOSETRON HYDROCHLORIDE, PALONOSETRON HYDROCHLORIDE
 PAMELOR, NORTRIPTYLINE HYDROCHLORIDE
 PAMIDRONATE DISODIUM, PAMIDRONATE DISODIUM
 PANCREAZE, PANCRELIPASE (AMYLASE)

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** P **

PANCURONIUM BROMIDE, PANCURONIUM BROMIDE
 PANDEL, HYDROCORTISONE PROBUTATE
 PANRETIN, ALITRETINOIN
 PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
 PARAGARD T 380A, COPPER
 PAREMYD, HYDROXYAMPHETAMINE HYDROBROMIDE
 PARICALCITOL, PARICALCITOL
 PARLODEL, BROMOCRIPTINE MESYLATE
 PARNATE, TRANYLCPROMINE SULFATE
 PAROEX, CHLORHEXIDINE GLUCONATE
 PAROMOMYCIN SULFATE, PAROMOMYCIN SULFATE
 PAROXETINE, PAROXETINE HYDROCHLORIDE
 PAROXETINE HYDROCHLORIDE, PAROXETINE HYDROCHLORIDE
 PAROXETINE MESYLATE, PAROXETINE MESYLATE
 PARSABIV, ETELCALCETIDE
 PASER, AMINOSALICYLIC ACID
 PATADAY, OLOPATADINE HYDROCHLORIDE
 PATANASE, OLOPATADINE HYDROCHLORIDE
 PATANOL, OLOPATADINE HYDROCHLORIDE
 PAXIL, PAROXETINE HYDROCHLORIDE
 PAXIL CR, PAROXETINE HYDROCHLORIDE
 PAZEO, OLOPATADINE HYDROCHLORIDE
 PEDIAPRED, PREDNISOLONE SODIUM PHOSPHATE
 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
 PEGANONE, ETHOTOIN
 PEMFEXY, PEMETREXED
 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
 PENLAC, CICLOPIROX
 PENNSAID, DICLOFENAC SODIUM
 PENTAM, PENTAMIDINE ISETHIONATE
 PENTAMIDINE ISETHIONATE, PENTAMIDINE ISETHIONATE
 PENTASA, MESALAMINE
 PENTETATE CALCIUM TRISODIUM, PENTETATE CALCIUM TRISODIUM
 PENTETATE ZINC TRISODIUM, PENTETATE ZINC TRISODIUM
 PENTOBARBITAL SODIUM, PENTOBARBITAL SODIUM
 PENTOLAIR, CYCLOPENTOLATE HYDROCHLORIDE
 PENTOSTATIN, PENTOSTATIN
 PENTOXIFYLLINE, PENTOXIFYLLINE
 PENTOXIL, 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
 PERIOCHIP, CHLORHEXIDINE GLUCONATE
 PERIOGARD, CHLORHEXIDINE GLUCONATE
 PERMETHRIN, PERMETHRIN (OTC)
 PERMETHRIN, PERMETHRIN
 PERPHENAZINE, PERPHENAZINE
 PERPHENAZINE AND AMITRIPTYLINE HYDROCHLORIDE, AMITRIPTYLINE HYDROCHLORIDE
 PERSANTINE, DIPYRIDAMOLE
 PERSERIS KIT, RISPERIDONE
 PERTZYE, PANCRELIPASE (AMYLASE)
 PEXEVA, PAROXETINE MESYLATE

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** P **

PFIZERPEN, PENICILLIN G POTASSIUM
 PHENDIMETRAZINE TARTRATE, PHENDIMETRAZINE TARTRATE
 PHENELZINE SULFATE, PHENELZINE SULFATE
 PHENOXYBENZAMINE HYDROCHLORIDE, PHENOXYBENZAMINE HYDROCHLORIDE
 PHENTERMINE HYDROCHLORIDE, 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
 PHILITH, ETHINYL ESTRADIOL
 PHOSLO GELCAPS, CALCIUM ACETATE
 PHOSLYRA, CALCIUM ACETATE
 PHOSPHOLINE IODIDE, ECHOTHIOPHATE IODIDE
 PHOTOFRIN, PORFIMER SODIUM
 PHOTREXA, RIBOFLAVIN 5'-PHOSPHATE SODIUM
 PHOTREXA VISCOUS IN DEXTRAN 20%, RIBOFLAVIN 5'-PHOSPHATE SODIUM
 PHOXILLUM B22K 4/0 IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 PHOXILLUM BK 4/2.5 IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 PHYSIOLYTE IN PLASTIC CONTAINER, MAGNESIUM CHLORIDE
 PHYSIOSOL IN PLASTIC CONTAINER, MAGNESIUM CHLORIDE
 PHYTONADIONE, PHYTONADIONE
 PICATO, INGENOL MEBUTATE
 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
 PIRMELLA 1/35, ETHINYL ESTRADIOL
 PIRMELLA 7/7/7, ETHINYL ESTRADIOL
 PIROXICAM, PIROXICAM
 PITAVASTATIN CALCIUM, PITAVASTATIN CALCIUM
 PITOCIN, OXYTOCIN
 PIZENSY, LACTITOL
 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
 PLAVIX, CLOPIDOGREL BISULFATE
 PLEGISOL IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 PLENVU, ASCORBIC ACID
 PLIAGLIS, LIDOCAINE
 PODOFILOX, PODOFILOX
 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
 POLYTRIM, POLYMYXIN B SULFATE
 POMALYST, POMALIDOMIDE
 PONSTEL, MEFENAMIC ACID
 PORTIA-28, ETHINYL ESTRADIOL
 POSACONAZOLE, POSACONAZOLE
 POTASSIUM ACETATE, POTASSIUM ACETATE
 POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
 POTASSIUM CHLORIDE 0.037% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER,

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** P **

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 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 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
 POVIDONE IODINE, POVIDONE-IODINE (OTC)
 PRADAXA, DABIGATRAN ETEXILATE MESYLATE
 PRALIDOXIME CHLORIDE, PRALIDOXIME CHLORIDE
 PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE
 PRAMOSONE, HYDROCORTISONE ACETATE
 PRASUGREL, PRASUGREL HYDROCHLORIDE
 PRAVACHOL, PRAVASTATIN SODIUM
 PRAVASTATIN SODIUM, PRAVASTATIN SODIUM
 PRAZICQUANTEL, PRAZICQUANTEL
 PRAZOSIN HYDROCHLORIDE, PRAZOSIN HYDROCHLORIDE
 PRE-OP, HEXACHLOROPHENE
 PRE-OP II, HEXACHLOROPHENE
 PRE-PEN, BENZYLPENICILLOYL POLYLYSINE
 PRECEDEX, DEXMEDETOMIDINE HYDROCHLORIDE
 PRED FORTE, PREDNISOLONE ACETATE
 PRED MILD, PREDNISOLONE ACETATE
 PRED-G, GENTAMICIN SULFATE
 PREDNICARBATE, PREDNICARBATE
 PREDNISOLONE, PREDNISOLONE
 PREDNISOLONE SODIUM PHOSPHATE, PREDNISOLONE SODIUM PHOSPHATE
 PREDNISONE, PREDNISONE
 PREDNISONE INTENSOL, PREDNISONE
 PREGABALIN, PREGABALIN
 PREGNYL, GONADOTROPIN, CHORIONIC
 PRELONE, PREDNISOLONE
 PREMARIN, ESTROGENS, CONJUGATED
 PREMASOL 10% IN PLASTIC CONTAINER, AMINO ACIDS
 PREMASOL 6% IN PLASTIC CONTAINER, AMINO ACIDS
 PREMYPHASE 14/14, ESTROGENS, CONJUGATED
 PREMPRO, ESTROGENS, CONJUGATED
 PREPIDIL, DINOPROSTONE
 PRESTALIA, AMLODIPINE BESYLATE
 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)
 PREVIFEM, ETHINYL ESTRADIOL
 PREVYMIS, LETERMOVIR
 PREZCOBIX, COBICISTAT
 PREZISTA, DARUNAVIR
 PRIALT, ZICONOTIDE ACETATE

APPENDIX A - PRODUCT NAME INDEX

** P **

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
 PRIMISOL, TRIMETHOPRIM HYDROCHLORIDE
 PRINIVIL, LISINAPRIL
 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 DIGIHALER, ALBUTEROL SULFATE
 PROAIR HFA, ALBUTEROL SULFATE
 PROAIR RESPICLICK, ALBUTEROL SULFATE
 PROBALAN, PROBENECID
 PROBENECID, PROBENECID
 PROBENECID AND COLCHICINE, COLCHICINE
 PROBUPHINE, BUPRENORPHINE HYDROCHLORIDE
 PROCAINAMIDE HYDROCHLORIDE, PROCAINAMIDE HYDROCHLORIDE
 PROCALAMINE, AMINO ACIDS
 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
 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
 PROMETH VC W/ CODEINE, CODEINE PHOSPHATE
 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 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
 PROPANTHELINE BROMIDE, PROPANTHELINE BROMIDE
 PROPARACAINE HYDROCHLORIDE, PROPARACAINE HYDROCHLORIDE
 PROPECIA, FINASTERIDE
 PROPOFOL, PROPOFOL
 PROPRANOLOL HYDROCHLORIDE, PROPRANOLOL HYDROCHLORIDE
 PROPRANOLOL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE

APPENDIX A - PRODUCT NAME INDEX**** P ****

PROPYLTHIOURACIL, PROPYLTHIOURACIL
PROSCAR, FINASTERIDE
PROSOL 20% SULFITE FREE IN PLASTIC CONTAINER, AMINO ACIDS
PROSTIN E2, DINOPROSTONE
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
PSEUDOEPHEDRINE HYDROCHLORIDE, PSEUDOEPHEDRINE HYDROCHLORIDE (OTC)
PULMICORT FLEXHALER, BUDESONIDE
PULMICORT RESPULES, BUDESONIDE
PUR-WASH, PURIFIED WATER (OTC)
PURINETHOL, MERCAPTOPURINE
PURIXAN, MERCAPTOPURINE
PYLERA, BISMUTH SUBCITRATE POTASSIUM
PYRAZINAMIDE, PYRAZINAMIDE
PYRIDOSTIGMINE BROMIDE, PYRIDOSTIGMINE BROMIDE
PYRIDOXINE HYDROCHLORIDE, PYRIDOXINE HYDROCHLORIDE
PYRIMETHAMINE, PYRIMETHAMINE
PYTEST, UREA, C-14
PYTEST KIT, UREA, C-14

**** Q ****

QBRELIS, LISINAPRIL
QBREXZA, GLYCOPYRROLIUM TOSYLATE
QMIIZ ODT, MELOXICAM
QNASL, BECLOMETHASONE DIPROPIONATE
QOLIANA, BRIMONIDINE TARTRATE
QSYMIA, PHENTERMINE HYDROCHLORIDE
QTERN, DAPAGLIFLOZIN
QTERNMET XR, DAPAGLIFLOZIN
QUADRAMET, SAMARIUM SM-153 LEXIDRONAM PENTASODIUM
QUALAQUIN, QUININE SULFATE
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
QUINARETIC, HYDROCHLOROTHIAZIDE
QUINIDINE GLUCONATE, QUINIDINE GLUCONATE
QUINIDINE SULFATE, QUINIDINE SULFATE
QUININE SULFATE, QUININE SULFATE
QUTENZA, CAPSAICIN
QUZYTIR, CETIRIZINE HYDROCHLORIDE
QVAR REDHALER, BECLOMETHASONE DIPROPIONATE

**** R ****

R-GENE 10, ARGININE HYDROCHLORIDE
RABEPRAZOLE SODIUM, RABEPRAZOLE SODIUM
RADICAVA, EDARAVONE
RADIOGARDASE (PRUSSIAN BLUE), FERRIC HEXACYANOFERRATE(II)
RADIOGENIX SYSTEM, TECHNETIUM TC-99M SODIUM PERTECHNETATE GENERATOR

APPENDIX A - PRODUCT NAME INDEX

** R **

RALOXIFENE HYDROCHLORIDE, RALOXIFENE HYDROCHLORIDE
 RAMELTEON, RAMELTEON
 RAMIPRIL, RAMIPRIL
 RANEXA, RANOLAZINE
 RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE (OTC)
 RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
 RANOLAZINE, RANOLAZINE
 RAPAFLO, SILODOSIN
 RAPAMUNE, SIROLIMUS
 RAPIVAB, PERAMIVIR
 RASAGILINE MESYLATE, RASAGILINE MESYLATE
 RASUVO, METHOTREXATE
 RAVICTI, GLYCEROL PHENYL BUTYRATE
 RAYALDEE, CALCIFEDIOL
 RAYOS, PREDNISONE
 RAZADYNE, GALANTAMINE HYDROBROMIDE
 RAZADYNE ER, GALANTAMINE HYDROBROMIDE
 READI-CAT 2, BARIUM SULFATE
 READI-CAT 2 SMOOTHIE, BARIUM SULFATE
 READYPREP CHG, CHLORHEXIDINE GLUCONATE (OTC)
 REBETOL, RIBAVIRIN
 RECARBRIO, CILASTATIN SODIUM
 RECLAST, ZOLEDRONIC ACID
 RECTIV, NITROGLYCERIN
 REDITREX, METHOTREXATE
 REGLAN, METOCLOPRAMIDE HYDROCHLORIDE
 REGONOL, PYRIDOSTIGMINE BROMIDE
 RELENZA, ZANAMIVIR
 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
 REPAGLINIDE, REPAGLINIDE
 REPRESXAIN, HYDROCODONE BITARTRATE
 REQUIP, ROPINIROLE HYDROCHLORIDE
 REQUIP XL, ROPINIROLE HYDROCHLORIDE
 RESCRIPTOR, DELAVIRDINE MESYLATE
 RESECTISOL IN PLASTIC CONTAINER, MANNITOL
 RESTASIS, CYCLOSPORINE
 RESTASIS MULTIDOSE, CYCLOSPORINE
 RESTORIL, TEMAZEPAM
 RETIN-A, TRETINOIN
 RETIN-A MICRO, TRETINOIN
 RETIN-A-MICRO, TRETINOIN
 RETISERT, FLUOCINOLONE ACETONIDE
 RETROVIR, ZIDOVUDINE
 REVATIO, SILDENAFIL CITRATE
 REVLIMID, LENALIDOMIDE
 REVONTO, DANTROLENE SODIUM
 REXULTI, BREXPIPIRAZOLE
 REYATAZ, ATAZANAVIR SULFATE
 REYVOW, LASMIDITAN SUCCINATE
 RHINOCORT ALLERGY, BUDESONIDE (OTC)
 RHOFAD, OXYMETAZOLINE HYDROCHLORIDE
 RHOPRESSA, NETARSUDIL MESYLATE
 RIBAVIRIN, RIBAVIRIN
 RIDAURA, AURANOFIN
 RIFABUTIN, RIFABUTIN
 RIFADIN, RIFAMPIN

APPENDIX A - PRODUCT NAME INDEX

** R **

RIFAMATE, ISONIAZID
 RIFAMPIN, RIFAMPIN
 RIFATER, ISONIAZID
 RILUTEK, RILUZOLE
 RILUZOLE, RILUZOLE
 RIMACTANE, RIFAMPIN
 RIMANTADINE HYDROCHLORIDE, RIMANTADINE HYDROCHLORIDE
 RIMSO-50, DIMETHYL SULFOXIDE
 RINGER'S IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 RINVOQ, UPADACITINIB
 RIOMET, METFORMIN HYDROCHLORIDE
 RIOMET ER, METFORMIN HYDROCHLORIDE
 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
 RIZATRIPTAN BENZOATE, RIZATRIPTAN BENZOATE
 ROBAXIN, METHOCARBAMOL
 ROBAXIN-750, METHOCARBAMOL
 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)
 ROPINIROLE HYDROCHLORIDE, ROPINIROLE HYDROCHLORIDE
 ROPIVACAINE HYDROCHLORIDE, ROPIVACAINE HYDROCHLORIDE
 ROSUVASTATIN CALCIUM, ROSUVASTATIN CALCIUM
 ROWASA, MESALAMINE
 ROXICET, ACETAMINOPHEN
 ROXICODONE, OXYCODONE HYDROCHLORIDE
 ROZEREM, RAMELTEON
 ROZLYTREK, ENTRECTINIB
 RUBRACA, RUCAPARIB CAMSYLATE
 RUBY-FILL, RUBIDIUM CHLORIDE RB-82
 RUFINAMIDE, RUFINAMIDE
 RUZURGI, AMIFAMPRIDINE
 RYANODEX, DANTROLENE SODIUM
 RYBELSUS, SEMAGLUTIDE
 RYDAPT, MIDOSTAURIN
 RYTARY, CARBIDOPA
 RYTHMOL SR, PROPAFENONE HYDROCHLORIDE

** S **

SABRIL, VIGABATRIN
 SAFYRAL, DROSPIRENONE
 SAIZEN, SOMATROPIN
 SALAGEN, PILOCARPINE HYDROCHLORIDE
 SALONPAS, MENTHOL (OTC)
 SAMSCA, TOLVAPTAN
 SANCUSO, GRANISETRON
 SANDIMMUNE, CYCLOSPORINE
 SANDOSTATIN, OCTREOTIDE ACETATE
 SANDOSTATIN LAR, OCTREOTIDE ACETATE
 SAPHRIS, ASENAFINE MALEATE
 SAPROPTERIN DIHYDROCHLORIDE, SAPROPTERIN DIHYDROCHLORIDE
 SARAFEM, FLUOXETINE HYDROCHLORIDE
 SAVAYSA, EDOXABAN TOSYLATE
 SAVELLA, MILNACIPRAN HYDROCHLORIDE

APPENDIX A - PRODUCT NAME INDEX

** S **

SAXENDA, LIRAGLUTIDE RECOMBINANT
 SCANDONEST L, LEVONORDEFIN
 SCANDONEST PLAIN, MEPIVACAINE HYDROCHLORIDE
 SCANLUX-300, IOPAMIDOL
 SCANLUX-370, IOPAMIDOL
 SCENESSE, AFAMELANOTIDE
 SCLEROSOL, TALC
 SCOPOLAMINE, SCOPOLAMINE
 SEASONALE, ETHINYL ESTRADIOL
 SEASONIQUE, ETHINYL ESTRADIOL
 SECONAL SODIUM, SECOBARBITAL SODIUM
 SECUADO, ASENAPINE
 SEEBRI, GLYCOPYRROLATE
 SEGLUROMET, ERTUGLIFLOZIN
 SEIZALAM, MIDAZOLAM HYDROCHLORIDE
 SELEGILINE HYDROCHLORIDE, SELEGILINE HYDROCHLORIDE
 SELENIUS ACID, SELENIUS ACID
 SELENIUM SULFIDE, SELENIUM SULFIDE
 SELFEMRA, FLUOXETINE HYDROCHLORIDE
 SELZENTRY, MARAVIROC
 SEMPREX-D, ACRIVASTINE
 SENSIPAR, CINACALCET HYDROCHLORIDE
 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
 SEROSTIM, SOMATROPIN
 SERTRALINE HYDROCHLORIDE, SERTRALINE HYDROCHLORIDE
 SETLAKIN, ETHINYL ESTRADIOL
 SEVELAMER CARBONATE, SEVELAMER CARBONATE
 SEVELAMER HYDROCHLORIDE, SEVELAMER HYDROCHLORIDE
 SEVOFLURANE, SEVOFLURANE
 SEYSARA, SARECYCLINE HYDROCHLORIDE
 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
 SIMPESE, ETHINYL ESTRADIOL
 SIMVASTATIN, SIMVASTATIN
 SINE-AID IB, IBUPROFEN (OTC)
 SINEMET, CARBIDOPA
 SINGULAIR, MONTELUKAST SODIUM
 SINUVA, MOMETASONE FUROATE
 SIROLIMUS, SIROLIMUS
 SIRTURO, BEDAQUILINE FUMARATE
 SITAVIG, ACYCLOVIR
 SIVEXTRO, TEDIZOLID PHOSPHATE
 SKELAXIN, METAXALONE
 SKLICE, IVERMECTIN
 SKYLA, LEVONORGESTREL
 SLYND, DROSPIRENONE
 SMOFLIPID 20%, FISH OIL
 SODIUM ACETATE, SODIUM ACETATE
 SODIUM BICARBONATE, SODIUM BICARBONATE
 SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, SODIUM CHLORIDE

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** S **

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 3% IN PLASTIC CONTAINER, SODIUM CHLORIDE
 SODIUM CHLORIDE 5% IN PLASTIC CONTAINER, SODIUM CHLORIDE
 SODIUM CHLORIDE IN PLASTIC CONTAINER, SODIUM CHLORIDE
 SODIUM FERRIC GLUCONATE COMPLEX IN SUCROSE, SODIUM FERRIC GLUCONATE COMPLEX
 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 LACTATE IN PLASTIC CONTAINER, SODIUM LACTATE
 SODIUM NITRITE, SODIUM NITRITE
 SODIUM NITROPRUSSIDE, SODIUM NITROPRUSSIDE
 SODIUM OXYBATE, SODIUM OXYBATE
 SODIUM PHENYLACETATE AND SODIUM BENZOATE, SODIUM BENZOATE
 SODIUM PHENYLBUTYRATE, SODIUM PHENYLBUTYRATE
 SODIUM PHOSPHATES IN PLASTIC CONTAINER, SODIUM PHOSPHATE, DIBASIC, HEPTAHYDRATE
 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
 SOJOURN, SEVOFLURANE
 SOLARAZE, DICLOFENAC SODIUM
 SOLIFENACIN SUCCINATE, SOLIFENACIN SUCCINATE
 SOLIQUA 100/33, INSULIN GLARGINE
 SOLODYN, MINOCYCLINE HYDROCHLORIDE
 SOLOSEC, SECNIDAZOLE
 SOLTAMOX, TAMOXIFEN CITRATE
 SOLU-CORTEF, HYDROCORTISONE SODIUM SUCCINATE
 SOLU-MEDROL, METHYLPREDNISOLONE SODIUM SUCCINATE
 SOLUPREP, CHLORHEXIDINE GLUCONATE (OTC)
 SOMA, CARISOPRODOL
 SOMATULINE DEPOT, LANREOTIDE ACETATE
 SOMAVERT, PEGVISOMANT
 SONATA, ZALEPLON
 SOOLANTRA, IVERMECTIN
 SORBITOL 3% IN PLASTIC CONTAINER, SORBITOL
 SORBITOL 3.3% IN PLASTIC CONTAINER, SORBITOL
 SORBITOL-MANNITOL IN PLASTIC CONTAINER, MANNITOL
 SORIATANE, ACITRETIN
 SORILUX, CALCIPOTRIENE
 SORINE, SOTALOL HYDROCHLORIDE
 SOTALOL HYDROCHLORIDE, SOTALOL HYDROCHLORIDE
 SOTRADECOL, SODIUM TETRADECYL SULFATE
 SOTYLIZE, SOTALOL HYDROCHLORIDE
 SOVALDI, SOFOSBUVIR
 SPECTAZOLE, ECONAZOLE NITRATE
 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

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** S **

STALEVO 50, CARBIDOPA
 STALEVO 75, CARBIDOPA
 STAVUDINE, STAVUDINE
 STAXYN, VARDENAFIL HYDROCHLORIDE
 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 IN PLASTIC CONTAINER, STERILE WATER FOR IRRIGATION
 STERITALC, TALC
 STIE-CORT, HYDROCORTISONE
 STIMATE (NEEDS NO REFRIGERATION), DESMOPRESSIN ACETATE
 STIOLTO RESPIMAT, OLODATEROL HYDROCHLORIDE
 STIVARGA, REGORAFENIB
 STRATTERA, ATOMOXETINE HYDROCHLORIDE
 STREPTOMYCIN SULFATE, STREPTOMYCIN SULFATE
 STRIBILD, COBICISTAT
 STRIVERDI RESPIMAT, OLODATEROL HYDROCHLORIDE
 STROMEKTOL, IVERMECTIN
 STRONTIUM CHLORIDE SR-89, STRONTIUM CHLORIDE SR-89
 SUBLIMAZE PRESERVATIVE FREE, FENTANYL CITRATE
 SUBLOCADE, BUPRENORPHINE
 SUBOXONE, BUPRENORPHINE HYDROCHLORIDE
 SUBSYS, FENTANYL
 SUCCINYLCHOLINE CHLORIDE, SUCCINYLCHOLINE CHLORIDE
 SUCRAID, SACROSIDASE
 SUCRALFATE, SUCRALFATE
 SUDAFED 12 HOUR, PSEUDOEPHEDRINE HYDROCHLORIDE (OTC)
 SUDAFED 24 HOUR, PSEUDOEPHEDRINE HYDROCHLORIDE (OTC)
 SUFENTA PRESERVATIVE FREE, SUFENTANIL CITRATE
 SUFENTANIL CITRATE, SUFENTANIL CITRATE
 SULAR, NISOLDIPINE
 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
 SUNOSI, SOLRIAMFETOL HYDROCHLORIDE
 SUPPRELIN LA, HISTRELIN ACETATE
 SUPRANE, DESFLURANE
 SUPRAX, CEFIXIME
 SUPREP BOWEL PREP KIT, MAGNESIUM SULFATE
 SURVANTA, BERACTANT
 SUSTIVA, EFAVIRENZ
 SUSTOL, GRANISETRON
 SUTENT, SUNITINIB MALATE
 SYEDA, DROSPIRENONE
 SYLEVIA, ETHINYL ESTRADIOL
 SYMBICORT, BUDESONIDE
 SYMBYAX, FLUOXETINE HYDROCHLORIDE
 SYMDEKO (COPACKAGED), IVACAFTOR
 SYMFI, EFAVIRENZ
 SYMFI LO, EFAVIRENZ
 SYMJEPI, EPINEPHRINE
 SYMLIN, PRAMLINTIDE ACETATE
 SYMPAZAN, CLOBAZAM
 SYMPROIC, NALDEMEDINE TOSYLATE

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** S **

SYMTUZA, COBICISTAT
 SYNALAR, FLUOCINOLONE ACETONIDE
 SYNAREL, NAFARELIN ACETATE
 SYNDROS, DRONABINOL
 SYNERA, LIDOCAINE
 SYNERCID, DALFOPRISTIN
 SYNJARDY, EMPAGLIFLOZIN
 SYNJARDY XR, EMPAGLIFLOZIN
 SYNRIPO, OMACETAXINE MEPESUCCINATE
 SYNTHROID, LEVOTHYROXINE SODIUM **
 SYPRINE, TRIENTINE HYDROCHLORIDE

** T **

TAB-PROFEN, IBUPROFEN (OTC)
 TACLONEX, BETAMETHASONE DIPROPIONATE
 TACROLIMUS, TACROLIMUS
 TADALAFIL, TADALAFIL
 TAFINLAR, DABRAFENIB MESYLATE
 TAFLUPROST, TAFLUPROST
 TAGAMET HB, CIMETIDINE (OTC)
 TAGITOL V, BARIUM SULFATE
 TAGRISSO, OSIMERTINIB MESYLATE
 TALC, TALC
 TALICIA, AMOXICILLIN
 TALZENNA, TALAZOPARIB TOSYLATE
 TAMBOCOR, FLECAINIDE ACETATE
 TAMIFLU, OSELTAMIVIR PHOSPHATE
 TAMOXIFEN CITRATE, TAMOXIFEN CITRATE
 TAMSULOSIN HYDROCHLORIDE, TAMSULOSIN HYDROCHLORIDE
 TAPAZOLE, METHIMAZOLE
 TARCEVA, ERLOTINIB HYDROCHLORIDE
 TARGRETIN, BEXAROTENE
 TARKA, TRANDOLAPRIL
 TASIGNA, NILOTINIB HYDROCHLORIDE
 TASMAR, TOLCAPONE
 TAVALISSE, FOSTAMATINIB DISODIUM
 TAXOTERE, DOCETAXEL
 TAYTULLA, ETHINYL ESTRADIOL
 TAZAROTENE, TAZAROTENE
 TAZICEF, CEFTAZIDIME
 TAZORAC, TAZAROTENE
 TAZTIA XT, DILTIAZEM HYDROCHLORIDE
 TAZVERIK, TAZEMETOSTAT HYDROBROMIDE
 TECFIDERA, DIMETHYL FUMARATE
 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 ALBUMIN AGGREGATED KIT, TECHNETIUM TC-99M ALBUMIN AGGREGATED KIT
 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
 TEFLARO, CEFTAROLINE FOSAMIL
 TEGRETOL, CARBAMAZEPINE
 TEGRETOL-XR, CARBAMAZEPINE
 TEGSEDI, INOTERSEN SODIUM
 TEKTURNA, ALISKIREN HEMIFUMARATE
 TEKTURNA HCT, ALISKIREN HEMIFUMARATE
 TELMISARTAN, TELMISARTAN
 TELMISARTAN AND AMLODIPINE, AMLODIPINE BESYLATE
 TELMISARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 TEMAZEPAM, TEMAZEPAM
 TEMIXYS, LAMIVUDINE
 TEMODAR, TEMOZOLOMIDE
 TEMOZOLOMIDE, TEMOZOLOMIDE

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** T **

TEMSIROLIMUS, TEMSIROLIMUS
 TENOFOVIR DISOPROXIL FUMARATE, TENOFOVIR DISOPROXIL FUMARATE
 TENORETIC 100, ATENOLOL
 TENORETIC 50, ATENOLOL
 TENORMIN, ATENOLOL
 TEPADINA, THIOTEPA
 TERAZOSIN HYDROCHLORIDE, TERAZOSIN HYDROCHLORIDE
 TERBINAFINE HYDROCHLORIDE, TERBINAFINE HYDROCHLORIDE (OTC)
 TERBINAFINE HYDROCHLORIDE, TERBINAFINE HYDROCHLORIDE
 TERBUTALINE SULFATE, TERBUTALINE SULFATE
 TERCONAZOLE, TERCONAZOLE
 TERIFLUNOMIDE, TERIFLUNOMIDE
 TERIL, CARBAMAZEPINE
 TESSALON, BENZONATATE
 TESTIM, TESTOSTERONE
 TESTOPEL, TESTOSTERONE
 TESTOSTERONE, TESTOSTERONE
 TESTOSTERONE CYPIONATE, TESTOSTERONE CYPIONATE
 TESTOSTERONE ENANTHATE, TESTOSTERONE ENANTHATE
 TESTRED, METHYLTESTOSTERONE
 TETRABENAZINE, TETRABENAZINE
 TETRACAINE HYDROCHLORIDE, TETRACAINE HYDROCHLORIDE
 TETRACYCLINE HYDROCHLORIDE, TETRACYCLINE HYDROCHLORIDE
 TEXACORT, HYDROCORTISONE
 THALLOUS CHLORIDE TL 201, THALLOUS CHLORIDE TL-201
 THALOMID, THALIDOMIDE
 THEO-24, THEOPHYLLINE
 THEOCHRON, THEOPHYLLINE
 THEOPHYLLINE, THEOPHYLLINE
 THEOPHYLLINE 0.04% AND DEXTROSE 5% IN PLASTIC CONTAINER, THEOPHYLLINE
 THEOPHYLLINE 0.08% AND DEXTROSE 5% IN PLASTIC CONTAINER, THEOPHYLLINE
 THEOPHYLLINE 0.16% AND DEXTROSE 5% IN PLASTIC CONTAINER, THEOPHYLLINE
 THEOPHYLLINE 0.32% AND DEXTROSE 5% IN PLASTIC CONTAINER, 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
 THYRO-TABS, LEVOTHYROXINE SODIUM **
 THYROGEN, THYROTROPIN ALFA
 THYROSAFE, POTASSIUM IODIDE (OTC)
 THYROSHIELD, 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)
 TIROSINT, LEVOTHYROXINE SODIUM
 TIROSINT-SOL, LEVOTHYROXINE SODIUM

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** T **

TIS-U-SOL, MAGNESIUM SULFATE
TIS-U-SOL IN PLASTIC CONTAINER, MAGNESIUM SULFATE
TISSUEBLUE, BRILLIANT BLUE G
TIVICAY, DOLUTEGRAVIR SODIUM
TIVORBEX, INDOMETHACIN
TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE
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
TOBRAMYCIN SULFATE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, TOBRAMYCIN SULFATE
TOBREX, TOBRAMYCIN
TODAY, NONOXYNOL-9 (OTC)
TOFRANIL, IMIPRAMINE HYDROCHLORIDE
TOLAK, FLUOROURACIL
TOLAZAMIDE, TOLAZAMIDE
TOLBUTAMIDE, TOLBUTAMIDE
TOLCAPONE, TOLCAPONE
TOLMETIN SODIUM, TOLMETIN SODIUM
TOLSURA, ITRACONAZOLE
TOLTERODINE TARTRATE, TOLTERODINE TARTRATE
TOPAMAX, TOPIRAMATE
TOPICORT, DESOXIMETASONE
TOPIRAMATE, TOPIRAMATE
TOPOTECAN HYDROCHLORIDE, TOPOTECAN HYDROCHLORIDE
TOPROL-XL, METOPROLOL SUCCINATE
TOREMIFENE CITRATE, TOREMIFENE CITRATE
TORISEL, TEMSIROLIMUS
TORSEMIDE, TORSEMIDE
TOSYMRA, SUMATRIPTAN
TOTECT, DEXRAZOXANE HYDROCHLORIDE
TOUJEO MAX SOLOSTAR, INSULIN GLARGINE RECOMBINANT
TOUJEO SOLOSTAR, INSULIN GLARGINE RECOMBINANT
TOVIAZ, FESOTERODINE FUMARATE
TPN ELECTROLYTES IN PLASTIC CONTAINER, CALCIUM CHLORIDE
TPOXX, TECOVIRIMAT
TRACLEER, BOSENTAN
TRADJENTA, LINAGLIPTIN
TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE
TRAMADOL HYDROCHLORIDE AND ACETAMINOPHEN, ACETAMINOPHEN
TRANDATE, LABETALOL HYDROCHLORIDE
TRANDOLAPRIL, TRANDOLAPRIL
TRANDOLAPRIL AND VERAPAMIL HYDROCHLORIDE, TRANDOLAPRIL
TRANEXAMIC ACID, TRANEXAMIC ACID
TRANSDERM SCOP, SCOPOLAMINE
TRANXENE, CLORAZEPATE DIPOTASSIUM
TRANLYCYPROMINE SULFATE, TRANLYCYPROMINE 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
TRESIBA, INSULIN DEGLUDEC
TRETINOIN, TRETINOIN
Trexall, METHOTREXATE SODIUM

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** T **

TREXIMET, NAPROXEN SODIUM
 TREZIX, ACETAMINOPHEN
 TRI LO SPRINTEC, ETHINYL ESTRADIOL
 TRI-ESTARYLLA, ETHINYL ESTRADIOL
 TRI-LEGEST 21, ETHINYL ESTRADIOL
 TRI-LEGEST FE, ETHINYL ESTRADIOL
 TRI-LINYAH, ETHINYL ESTRADIOL
 TRI-LO-ESTARYLLA, ETHINYL ESTRADIOL
 TRI-LO-MILI, ETHINYL ESTRADIOL
 TRI-LUMA, FLUOCINOLONE ACETONIDE
 TRI-MILI, ETHINYL ESTRADIOL
 TRI-NORINYL 28-DAY, ETHINYL ESTRADIOL
 TRI-PREVIFEM, ETHINYL ESTRADIOL
 TRI-SPRINTEC, ETHINYL ESTRADIOL
 TRIACIN-C, CODEINE PHOSPHATE
 TRIAMCINOLONE ACETATE, TRIAMCINOLONE ACETONIDE
 TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE (OTC)
 TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE
 TRIAMCINOLONE ACETONIDE IN ABSORBASE, TRIAMCINOLONE ACETONIDE
 TRIAMTERENE, TRIAMTERENE
 TRIAMTERENE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 TRIAZOLAM, TRIAZOLAM
 TRIBENZOR, AMLODIPINE BESYLATE
 TRICOR, FENOFIBRATE
 TRIDERM, TRIAMCINOLONE ACETONIDE
 TRIENTINE HYDROCHLORIDE, TRIENTINE HYDROCHLORIDE
 TRISENCE, TRIAMCINOLONE ACETONIDE
 TRIFERIC, FERRIC PYROPHOSPHATE CITRATE
 TRIFLUOPERAZINE HYDROCHLORIDE, TRIFLUOPERAZINE HYDROCHLORIDE
 TRIFLURIDINE, TRIFLURIDINE
 TRIGLIDE, FENOFIBRATE
 TRIHEXYPHENIDYL HYDROCHLORIDE, TRIHEXYPHENIDYL HYDROCHLORIDE
 TRIJARDY XR, EMPAGLIFLOZIN
 TRIKAFTA (COPACKAGED), ELEXACAF TOR, IVACAFTOR, TEZACAFTOR
 TRILEPTAL, OXCARBAZEPINE
 TRILIPIX, CHOLINE FENOFIBRATE
 TRILYTE, POLYETHYLENE GLYCOL 3350
 TRIMETHOBENZAMIDE HYDROCHLORIDE, TRIMETHOBENZAMIDE HYDROCHLORIDE
 TRIMETHOPRIM, TRIMETHOPRIM
 TRIMETHOPRIM SULFATE AND POLYMYXIN B SULFATE, POLYMYXIN B SULFATE
 TRIMIPRAMINE MALEATE, TRIMIPRAMINE MALEATE
 TRINTELLIX, VORTIOXETINE HYDROBROMIDE
 TRIOSTAT, LIOTHYRONINE SODIUM
 TRIPTODUR KIT, TRIPTORELIN PAMOATE
 TRISENOX, ARSENIC TRIOXIDE
 TRIUMEQ, ABACAVIR SULFATE
 TRIVAGIZOLE 3, CLOTRIMAZOLE (OTC)
 TRIVORA-28, ETHINYL ESTRADIOL
 TRIZIVIR, ABACAVIR SULFATE
 TROKENDI XR, TOPIRAMATE
 TROPHAMINE, AMINO ACIDS
 TROPHAMINE 10%, AMINO ACIDS
 TROPICACYL, TROPICAMIDE
 TROPICAMIDE, TROPICAMIDE
 TROSPIUM CHLORIDE, TROSPIUM CHLORIDE
 TRULANCE, PLECANATIDE
 TRUSOPT, DORZOLAMIDE HYDROCHLORIDE
 TRUVADA, EMTRICITABINE
 TUDORZA PRESSAIR, ACLIDINIUM BROMIDE
 TURALIO, PEXIDARTINIB HYDROCHLORIDE
 TUSSICAPS, CHLORPHENIRAMINE POLISTIREX
 TUXARIN ER, CHLORPHENIRAMINE MALEATE
 TUZISTRA XR, CHLORPHENIRAMINE POLISTIREX
 TWIRLA, ETHINYL ESTRADIOL
 TWYNSTA, AMLODIPINE BESYLATE

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** T **

TYBOST, COBICISTAT
 TYDEMY, DROSPIRENONE
 TYGACIL, TIGECYCLINE
 TYKERB, LAPATINIB DITOSYLATE
 TYLENOL, ACETAMINOPHEN (OTC)
 TYLENOL W/ CODEINE NO. 3, ACETAMINOPHEN
 TYLENOL W/ CODEINE NO. 4, ACETAMINOPHEN
 TYMLOS, ABALOPARATIDE
 TYVASO, TREPROSTINIL
 TYZINE, TETRAHYDROZOLINE HYDROCHLORIDE

** U **

U-CORT, HYDROCORTISONE ACETATE
 UBRELVY, UBROGEPANT
 UCERIS, BUDESONIDE
 ULORIC, FEBUXOSTAT
 ULTACAN, ARTICAINE HYDROCHLORIDE
 ULTANE, SEVOFLURANE
 ULTIVA, REMIFENTANIL HYDROCHLORIDE
 ULTRA-TECHNEKOW FM, TECHNETIUM TC-99M SODIUM PERTECHNETATE GENERATOR
 ULTRACET, ACETAMINOPHEN
 ULTRAM, TRAMADOL HYDROCHLORIDE
 ULTRATAG, TECHNETIUM TC-99M RED BLOOD CELL KIT
 ULTRAVATE, HALOBETASOL PROPIONATE
 ULTRAVIST (PHARMACY BULK), IOPROMIDE
 ULTRAVIST 300, IOPROMIDE
 ULTRAVIST 370, IOPROMIDE
 UNASYN, AMPICILLIN SODIUM
 UNISOM, DOXYLAMINE SUCCINATE (OTC)
 UNITHROID, LEVOTHYROXINE SODIUM **
 UPTRAVI, SELEXIPAG
 URECHOLINE, BETHANECHOL CHLORIDE
 UREX, METHENAMINE HIPPURATE
 UROCIT-K, POTASSIUM CITRATE
 UROXATRAL, ALFUZOSIN HYDROCHLORIDE
 URSO 250, URSODIOL
 URSO FORTE, URSODIOL
 URSODIOL, URSODIOL
 UTIBRON, GLYCOPYRROLATE
 UVADEX, METHOXSALEN

** V **

VABOMERE, MEROPENEM
 VAGIFEM, ESTRADIOL
 VAGISTAT-1, TIOCONAZOLE (OTC)
 VALACYCLOVIR HYDROCHLORIDE, VALACYCLOVIR HYDROCHLORIDE
 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
 VANCOCIN HYDROCHLORIDE IN PLASTIC CONTAINER, VANCOMYCIN HYDROCHLORIDE
 VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE
 VANCOMYCIN HYDROCHLORIDE IN PLASTIC CONTAINER, VANCOMYCIN HYDROCHLORIDE
 VANDAZOLE, METRONIDAZOLE
 VANIQA, EFLORNITHINE HYDROCHLORIDE

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** v **

VANOS, FLUOCINONIDE
 VANTAS, HISTRELIN ACETATE
 VAPRISOL IN 5% DEXTROSE IN PLASTIC CONTAINER, CONIVAPTAN HYDROCHLORIDE
 VARDENAFIL HYDROCHLORIDE, VARDENAFIL HYDROCHLORIDE
 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
 VASOSTRICT, VASOPRESSIN
 VASOTEC, ENALAPRIL MALEATE
 VAZALORE, ASPIRIN (OTC)
 VAZCULEP, PHENYLEPHRINE HYDROCHLORIDE
 VECTICAL, CALCITRIOL
 VECURONIUM BROMIDE, VECURONIUM BROMIDE
 VELCADE, BORTEZOMIB
 VELETRI, EPOPROSTENOL SODIUM
 VELIVET, DESOGESTREL
 VELPHORO, SUCROFERRIC OXYHYDROXIDE
 VELTASSA, PATIROMER SORBITEX CALCIUM
 VELTIN, CLINDAMYCIN PHOSPHATE
 VEMLIDY, TENOFOVIR ALAFENAMIDE FUMARATE
 VENCLEXTA, VENETOCLAX
 VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE
 VENOFER, IRON SUCROSE
 VENTAVIS, ILOPROST
 VENTOLIN HFA, ALBUTEROL SULFATE
 VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE
 VERDESO, DESONIDE
 VEREGEN, SINECATECHINS
 VERELAN, VERAPAMIL HYDROCHLORIDE
 VERELAN PM, VERAPAMIL HYDROCHLORIDE
 VERSACLOZ, CLOZAPINE
 VERZENIO, ABEMACICLIB
 VESICARE, SOLIFENACIN SUCCINATE
 VFEND, VORICONAZOLE
 VIAGRA, SILDENAFIL CITRATE
 VIBATIV, TELAVANCIN HYDROCHLORIDE
 VIBERZI, ELUXADOLINE
 VIBISONE, CYANOCOBALAMIN
 VIBRAMYCIN, DOXYCYCLINE
 VIBRAMYCIN, DOXYCYCLINE CALCIUM
 VIBRAMYCIN, DOXYCYCLINE HYCLATE
 VICTOZA, LIRAGLUTIDE RECOMBINANT
 VIDAZA, AZACITIDINE
 VIDEX, DIDANOSINE
 VIDEX EC, DIDANOSINE
 VIEKIRA PAK (COPACKAGED), DASABUVIR SODIUM
 VIENVA, ETHINYL ESTRADIOL
 VIGABATRIN, VIGABATRIN
 VIGADRONE, VIGABATRIN
 VIGAMOX, MOXIFLOXACIN HYDROCHLORIDE
 VIIBRYD, VILAZODONE HYDROCHLORIDE
 VILAZODONE HYDROCHLORIDE, VILAZODONE HYDROCHLORIDE
 VIMOVO, ESOMEPRAZOLE MAGNESIUM
 VIMPAT, LACOSAMIDE
 VINBLASTINE SULFATE, VINBLASTINE SULFATE
 VINCRISTINE SULFATE PFS, VINCRISTINE SULFATE
 VINORELBINE TARTRATE, VINORELBINE TARTRATE
 VIOKACE, PANCRELIPASE (AMYLASE)
 VIORELE, DESOGESTREL

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** V **

VIRACEPT, NELFINAVIR MESYLATE
 VIRAMUNE, NEVIRAPINE
 VIRAMUNE XR, NEVIRAPINE
 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
 VISTOGARD, URIDINE TRIACETATE
 VISUDYNE, VERTEPORFIN
 VITAMIN D, ERGOCALCIFEROL
 VITAMIN K1, PHYTONADIONE
 VITRAKVI, LAROTRECTINIB SULFATE
 VITRASE, HYALURONIDASE
 VIVACTIL, PROTRIPTYLINE HYDROCHLORIDE
 VIVELLE-DOT, ESTRADIOL
 VIVITROL, NALTREXONE
 VIVLODEX, MELOXICAM
 VIZAMYL, FLUTEMETAMOL F-18
 VIZIMPRO, DACOMITINIB
 VOGELXO, TESTOSTERONE
 VOLNEA, DESOGESTREL
 VOLTAREN, DICLOFENAC SODIUM
 VORICONAZOLE, VORICONAZOLE
 VOSEVI, SOFOSBUVIR
 VOSOL, ACETIC ACID, GLACIAL
 VOSOL HC, ACETIC ACID, GLACIAL
 VOSPIRE ER, ALBUTEROL SULFATE
 VOTRIENT, PAZOPANIB HYDROCHLORIDE
 VPRIV, VELAGLUCERASE ALFA
 VRAYLAR, CARIPRAZINE HYDROCHLORIDE
 VUMERITY, DIROXIMEL FUMARATE
 VUSION, MICONAZOLE NITRATE
 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 **

WAKIX, PITOLISANT HYDROCHLORIDE
 WARFARIN SODIUM, WARFARIN SODIUM
 WELCHOL, COLESEVELAM HYDROCHLORIDE
 WELLBUTRIN SR, BUPROPION HYDROCHLORIDE
 WELLBUTRIN XL, BUPROPION HYDROCHLORIDE
 WERA, ETHINYL ESTRADIOL
 WIXELA INHUB, FLUTICASONE PROPIONATE
 WOMEN'S ROGAINE, MINOXIDIL (OTC)

** X **

XADAGO, SAFINAMIDE MESYLATE
 XALATAN, LATANOPROST
 XALKORI, CRIZOTINIB
 XANAX, ALPRAZOLAM
 XANAX XR, ALPRAZOLAM
 XARELTO, RIVAROXABAN
 XATMEP, METHOTREXATE SODIUM
 XELJANZ, TOFACITINIB CITRATE

APPENDIX A - PRODUCT NAME INDEX

** X **

XELJANZ XR, TOFACITINIB CITRATE
 XELODA, CAPECITABINE
 XELPROS, LATANOPROST
 XENAZINE, TETRABENAZINE
 XENICAL, ORLISTAT
 XENLETA, LEFAMULIN ACETATE
 XENON XE 133, XENON XE-133
 XEPI, OZENOXACIN
 XERAVA, ERAVACYCLINE DIHYDROCHLORIDE
 XERESE, ACYCLOVIR
 XERMELO, TELOTRISTAT ETIPRATE
 XHANCE, FLUTICASONE PROPIONATE
 XIFAXAN, RIFAXIMIN
 XIGDUO XR, DAPAGLIFLOZIN
 XIIDRA, LIFITEGRAST
 XIMINO, MINOCYCLINE HYDROCHLORIDE
 XOFIGO, RADIUM RA-223 DICHLORIDE
 XOFLUZA, BALOXAVIR MARBOXIL
 XOLEGEL, KETOCONAZOLE
 XOPENEX, LEVALBUTEROL HYDROCHLORIDE
 XOPENEX HFA, LEVALBUTEROL TARTRATE
 XOSPATA, GILTERITINIB FUMARATE
 XPOVIO, SELINEXOR
 XTAMPZA ER, OXYCODONE
 XTANDI, ENZALUTAMIDE
 XULANE, ETHINYL ESTRADIOL
 XULTOPHY 100/3.6, INSULIN DEGLUDEC
 XURIDEN, URIDINE TRIACETATE
 XYLOCAINE, LIDOCAINE HYDROCHLORIDE
 XYLOCAINE W/ EPINEPHRINE, EPINEPHRINE
 XYOSTED (AUTOINJECTOR), TESTOSTERONE ENANTHATE
 XYREM, SODIUM OXYBATE
 XYZAL, LEVOCETIRIZINE DIHYDROCHLORIDE
 XYZAL ALLERGY 24HR, LEVOCETIRIZINE DIHYDROCHLORIDE (OTC)

** Y **

YAELA, DROSPIRENONE
 YASMIN, DROSPIRENONE
 YAZ, DROSPIRENONE
 YONDELIS, TRABECTEDIN
 YONSA, ABIRATERONE ACETATE
 YOSPRALA, ASPIRIN
 YUPELRI, REVEFENACIN
 YUTIQ, FLUOCINOLONE ACETONIDE

** Z **

ZADITOR, KETOTIFEN FUMARATE (OTC)
 ZAFIRLUKAST, ZAFIRLUKAST
 ZALEPLON, ZALEPLON
 ZANAFLEX, TIZANIDINE HYDROCHLORIDE
 ZANOSAR, STREPTOZOCIN
 ZANTAC, RANITIDINE HYDROCHLORIDE
 ZANTAC 150, RANITIDINE HYDROCHLORIDE (OTC)
 ZANTAC 75, RANITIDINE HYDROCHLORIDE (OTC)
 ZARONTIN, ETHOSUXIMIDE
 ZAROXOLYN, METOLAZONE
 ZAVESCA, MIGLUSTAT
 ZEGERID, OMEPRAZOLE
 ZEGERID OTC, OMEPRAZOLE (OTC)
 ZEJULA, NIRAPARIB TOSYLATE
 ZELAPAR, SELEGILINE HYDROCHLORIDE
 ZELBORAF, VEMURAFENIB
 ZELNORM, TEGASEROD MALEATE
 ZEMBRACE SYMTOUCH, SUMATRIPTAN SUCCINATE
 ZEMDRI, PLAZOMICIN SULFATE

APPENDIX A - PRODUCT NAME INDEX

** Z **

ZEMPLAR, PARICALCITOL
ZENATANE, ISOTRETINOIN
ZENPEP, PANCRELIPASE (AMYLASE)
ZEPATIER, ELBASVIR
ZERBAXA, CEFTOLOZANE SULFATE
ZERIT, STAVUDINE
ZERVIAE, CETIRIZINE HYDROCHLORIDE
ZESTORETIC, HYDROCHLOROTHIAZIDE
ZESTRIL, LISINAPRIL
ZETIA, EZETIMIBE
ZETONNA, CICLESONIDE
ZIAC, BISOPROLOL FUMARATE
ZIAGEN, ABACAVIR SULFATE
ZIANA, CLINDAMYCIN PHOSPHATE
ZIDOVUDINE, ZIDOVUDINE
ZILEUTON, ZILEUTON
ZILRETTA, TRIAMCINOLONE ACETONIDE
ZINACEF, CEFUROXIME SODIUM
ZINC CHLORIDE IN PLASTIC CONTAINER, ZINC CHLORIDE
ZINC SULFATE, ZINC SULFATE
ZINECARD, DEXRAZOXANE HYDROCHLORIDE
ZINGO, LIDOCAINE HYDROCHLORIDE
ZIOPTAN, TAFLUPROST
ZIPRASIDONE HYDROCHLORIDE, ZIPRASIDONE HYDROCHLORIDE
ZIPRASIDONE MESYLATE, ZIPRASIDONE MESYLATE
ZIPSOR, DICLOFENAC POTASSIUM
ZIRGAN, GANCICLOVIR
ZITHROMAX, AZITHROMYCIN
ZOCOR, SIMVASTATIN
ZOFRAN, ONDANSETRON HYDROCHLORIDE
ZOFRAN ODT, ONDANSETRON
ZOHYDRO ER, HYDROCODONE BITARTRATE
ZOLADEX, GOSERELIN ACETATE
ZOLEDRONIC, ZOLEDRONIC ACID
ZOLEDRONIC ACID, ZOLEDRONIC ACID
ZOLINZA, VORINOSTAT
ZOLMITRIPTAN, ZOLMITRIPTAN
ZOLOFT, SERTRALINE HYDROCHLORIDE
ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE
ZOLPIMIST, ZOLPIDEM TARTRATE
ZOMACTON, SOMATROPIN
ZOMETA, ZOLEDRONIC ACID
ZOMIG, ZOLMITRIPTAN
ZOMIG-ZMT, ZOLMITRIPTAN
ZONALON, DOXEPIN HYDROCHLORIDE
ZONEGRAN, ZONISAMIDE
ZONISAMIDE, ZONISAMIDE
ZONTIVITY, VORAPAXAR SULFATE
ZORBTIVE, SOMATROPIN RECOMBINANT
ZORTRESS, EVEROLIMUS
ZORVOLEX, DICLOFENAC
ZOSYN, PIPERACILLIN SODIUM
ZOSYN IN PLASTIC CONTAINER, PIPERACILLIN SODIUM
ZOVIA 1/35E-28, ETHINYL ESTRADIOL
ZOVIA 1/50E-28, ETHINYL ESTRADIOL
ZOVIRAX, ACYCLOVIR
ZTLIDO, LIDOCAINE
ZUBSOLV, BUPRENORPHINE HYDROCHLORIDE
ZULRESSO, BREXANOLONE
ZUMANDIMINE, DROSPIRENONE
ZUPLENZ, ONDANSETRON
ZYCLARA, IMIQUIMOD
ZYDELIG, IDELALISIB
ZYFLO, ZILEUTON
ZYFLO CR, ZILEUTON

APPENDIX A - PRODUCT NAME INDEX

**** Z ****

ZYKADIA, CERITINIB
ZYLET, LOTEPREDNOL ETABONATE
ZYLOPRIM, ALLOPURINOL
ZYMAR, GATIFLOXACIN
ZYMAXID, GATIFLOXACIN
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 DRUG DELIVERY

- * 3M DRUG DELIVERY SYSTEMS
 FENTANYL-100, FENTANYL
 FENTANYL-12, FENTANYL
 FENTANYL-25, FENTANYL
 FENTANYL-50, FENTANYL
 FENTANYL-75, FENTANYL
 PROVENTIL-HFA, ALBUTEROL SULFATE

3M HEALTH CARE

- * 3M HEALTH CARE INFECTION PREVENTION DIV
 SOLUPREP, CHLORHEXIDINE GLUCONATE (OTC)

**** 6 ****

60 DEGREES PHARMS

- * 60 DEGREES PHARMACEUTICALS LLC
 ARAKODA, TAFENOQUINE SUCCINATE

**** A ****

AAA USA INC

- * ADVANCED ACCELERATOR APPLICATIONS USA INC
 LUTATHERA, LUTETIUM DOTATATE LU-177
 NETSPOT, GALLIUM DOTATATE GA-68

AAIPHARMA LLC

- * AAIPHARMA LLC
 AZASAN, AZATHIOPRINE

ABBVIE

- * ABBVIE INC
 ANDROGEL, TESTOSTERONE
 CREON, PANCRELIPASE (AMYLASE)
 CYCLOSPORINE, CYCLOSPORINE
 DEPAKOTE ER, DIVALPROEX SODIUM
 DEPAKOTE, DIVALPROEX SODIUM
 GENGRAF, CYCLOSPORINE
 K-TAB, POTASSIUM CHLORIDE
 KALETRA, LOPINAVIR
 NIASPAN, NIACIN
 NIMBEX PRESERVATIVE FREE, CISATRACURIUM BESYLATE
 NIMBEX, CISATRACURIUM BESYLATE
 NORVIR, RITONAVIR
 SURVANTA, BERACTANT
 SYNTHROID, LEVOTHYROXINE SODIUM **
 TARKA, TRANDOLAPRIL
 TRICOR, FENOFIBRATE
 TRILIPIX, CHOLINE FENOFIBRATE
 ULTANE, SEVOFLURANE
 ZEMPLAR, PARICALCITOL

ABBVIE ENDOCRINE

- * ABBVIE ENDOCRINE INC
 LUPANETA PACK, LEUPROLIDE ACETATE

ABBVIE ENDOCRINE INC

- * ABBVIE ENDOCRINE INC
 LUPRON DEPOT, LEUPROLIDE ACETATE
 LUPRON DEPOT-PED, LEUPROLIDE ACETATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

**** A ****

ABBVIE INC

* ABBVIE INC
 DUOPA, CARBIDOPA
 MAVYRET, GLECAPREVIR
 NORVIR, RITONAVIR
 ORILISSA, ELAGOLIX SODIUM
 RINVOQ, UPADACITINIB
 VENCLEXTA, VENETOCLAX
 VIEKIRA PAK (COPACKAGED), DASABUVIR SODIUM

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
 CLOFARABINE, CLOFARABINE

ABRAXIS BIOSCIENCE

* ABRAXIS BIOSCIENCE LLC
 ABRAXANE, PACLITAXEL

ABRAXIS PHARM

* ABRAXIS PHARMACEUTICAL PRODUCTS
 CLINDAMYCIN PHOSPHATE IN DEXTROSE 5%, CLINDAMYCIN PHOSPHATE

ACACIA PHARMA LTD

* ACACIA PHARMA LTD
 BARHEMSYS, AMISULPRIDE

ACADIA PHARMS INC

* ACADIA PHARMACEUTICALS INC
 NUPLAZID, PIMAVANSERIN TARTRATE

ACCELRX LABS

* ACCELRX LABS LLC
 CARISOPRODOL, CARISOPRODOL

ACCORD HLTHCARE

* ACCORD HEALTHCARE INC
 ALLOPURINOL, ALLOPURINOL
 AMITRIPTYLINE HYDROCHLORIDE, AMITRIPTYLINE HYDROCHLORIDE
 AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
 ANASTROZOLE, ANASTROZOLE
 ARIPIPIRAZOLE, ARIPIPIRAZOLE
 ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM
 AZACITIDINE, AZACITIDINE
 BICALUTAMIDE, BICALUTAMIDE
 BIVALIRUDIN, BIVALIRUDIN
 BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
 BUSPIRONE HYDROCHLORIDE, BUSPIRONE HYDROCHLORIDE
 CAPECITABINE, CAPECITABINE
 CARBIDOPA AND LEVODOPA, CARBIDOPA
 CARBOPLATIN, CARBOPLATIN
 CISPLATIN, CISPLATIN
 CLOFARABINE, CLOFARABINE
 CLONAZEPAM, CLONAZEPAM
 CLOPIDOGREL BISULFATE, CLOPIDOGREL BISULFATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

**** A ****

* ACCORD HEALTHCARE INC
 CLOZAPINE, CLOZAPINE
 DALFAMPRIDINE, DALFAMPRIDINE
 DAPTOMYCIN, DAPTOMYCIN
 DECITABINE, DECITABINE
 DESOGESTREL AND ETHINYL ESTRADIOL, DESOGESTREL
 DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE
 DOCETAXEL, DOCETAXEL
 DOXAZOSIN MESYLATE, DOXAZOSIN MESYLATE
 DROSPIRENONE AND ETHINYL ESTRADIOL, DROSPIRENONE
 ENTECAVIR, ENTECAVIR
 EPLERENONE, EPLERENONE
 EPTIFIBATIDE, EPTIFIBATIDE
 ESCITALOPRAM OXALATE, ESCITALOPRAM OXALATE
 ESOMEPRAZOLE SODIUM, ESOMEPRAZOLE SODIUM
 ETOPOSIDE, ETOPOSIDE
 EZETIMIBE, EZETIMIBE
 FINASTERIDE, FINASTERIDE
 FLUOROURACIL, FLUOROURACIL
 GEMCITABINE HYDROCHLORIDE, GEMCITABINE HYDROCHLORIDE
 GLIMEPIRIDE, GLIMEPIRIDE
 GLIPIZIDE, GLIPIZIDE
 HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 IBANDRONATE SODIUM, IBANDRONATE SODIUM
 IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE
 ITRACONAZOLE, ITRACONAZOLE
 LETROZOLE, LETROZOLE
 LEVETIRACETAM, LEVETIRACETAM
 LISINOPRIL, LISINOPRIL
 LURASIDONE HYDROCHLORIDE, LURASIDONE HYDROCHLORIDE
 METHOTREXATE SODIUM PRESERVATIVE FREE, METHOTREXATE SODIUM
 METHOTREXATE SODIUM, METHOTREXATE SODIUM
 METHYLDOPA, METHYLDOPA
 MITOMYCIN, MITOMYCIN
 MONTELUKAST SODIUM, MONTELUKAST SODIUM
 MYCOPHENOLATE MOFETIL, MYCOPHENOLATE MOFETIL
 MYCOPHENOLIC ACID, MYCOPHENOLIC ACID
 NALTREXONE HYDROCHLORIDE, NALTREXONE HYDROCHLORIDE
 NORETHINDRONE AND ETHINYL ESTRADIOL AND FERROUS FUMARATE, ETHINYL ESTRADIOL
 NORETHINDRONE AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
 NORETHINDRONE, NORETHINDRONE
 OLMESARTAN MEDOXOMIL, OLMESARTAN MEDOXOMIL
 ONDANSETRON HYDROCHLORIDE PRESERVATIVE FREE, ONDANSETRON HYDROCHLORIDE
 ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
 OXALIPLATIN, OXALIPLATIN
 OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE
 PARICALCITOL, PARICALCITOL
 PIOGLITAZONE HYDROCHLORIDE, PIOGLITAZONE HYDROCHLORIDE
 PRASUGREL, PRASUGREL HYDROCHLORIDE
 PRAVASTATIN SODIUM, PRAVASTATIN SODIUM
 QUETIAPINE FUMARATE, QUETIAPINE FUMARATE
 RAMIPRIL, RAMIPRIL
 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

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

**** A ****

* ACCORD HEALTHCARE INC
 ZOLEDRONIC ACID, ZOLEDRONIC ACID

ACCORD HLTHCARE INC

* ACCORD HEALTHCARE INC USA
 BUSULFAN, BUSULFAN
 TIGECYCLINE, TIGECYCLINE

ACELLA

* ACELLA PHARMACEUTICALS LLC
 BROMPHENIRAMINE MALEATE, PSEUDOEPHEDRINE HYDROCHLORIDE AND DEXTROMETHORPHAN
 CICLOPIROX, CICLOPIROX
 PHENYTOIN SODIUM, PHENYTOIN SODIUM

ACELLA PHARMS LLC

* ACELLA PHARMACEUTICALS LLC
 GABAPENTIN, GABAPENTIN

ACELRX PHARMS

* ACELRX PHARMACEUTICALS INC
 DSUVIA, SUFENTANIL CITRATE

ACERUS

* ACERUS PHARMACEUTICALS CORP
 NATESTO, TESTOSTERONE

ACI HEALTHCARE LTD

* ACI HEALTHCARE LTD
 DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
 GABAPENTIN, GABAPENTIN
 LEVETIRACETAM, LEVETIRACETAM
 TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE

ACIC PHARMS

* ACIC PHARMACEUTICALS INC
 LEVETIRACETAM, LEVETIRACETAM
 RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
 TRANEXAMIC ACID, TRANEXAMIC ACID

ACORDA

* ACORDA THERAPEUTICS INC
 AMPYRA, DALFAMPRIDINE
 INBRIJA, LEVODOPA

ACP NIMBLE

* ACP NIMBLE BUYER INC
 ACEPHEN, ACETAMINOPHEN (OTC)
 ACYCLOVIR, ACYCLOVIR
 BETA-VAL, BETAMETHASONE VALERATE
 BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
 CALCIPOTRIENE, CALCIPOTRIENE
 CICLOPIROX, CICLOPIROX
 CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE
 CLINDAMYCIN HYDROCHLORIDE, CLINDAMYCIN HYDROCHLORIDE
 CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
 DESONIDE, DESONIDE
 DESOXIMETASONE, DESOXIMETASONE
 DOXYCYCLINE, DOXYCYCLINE
 ENALAPRIL MALEATE AND HYDROCHLOROTHIAZIDE, ENALAPRIL MALEATE
 FLUOCINOLONE ACETONIDE, FLUOCINOLONE ACETONIDE
 FLUOCINONIDE EMULSIFIED BASE, FLUOCINONIDE
 FLUOCINONIDE, FLUOCINONIDE
 FLUTICASONE PROPIONATE, FLUTICASONE PROPIONATE
 GENTAMICIN SULFATE, GENTAMICIN SULFATE
 HALOBETASOL PROPIONATE, HALOBETASOL PROPIONATE
 INDOMETHACIN, INDOMETHACIN
 LIDOCAINE, LIDOCAINE
 METRONIDAZOLE, METRONIDAZOLE
 MICONAZOLE NITRATE, MICONAZOLE NITRATE (OTC)
 MOMETASONE FUROATE, MOMETASONE FUROATE
 MYKACET, NYSTATIN
 NYSTATIN, NYSTATIN
 PROCHLORPERAZINE, PROCHLORPERAZINE
 PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

**** A ****

* ACP NIMBLE BUYER INC
 PROMETHEGAN, PROMETHAZINE HYDROCHLORIDE
 TAZAROTENE, TAZAROTENE
 TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE

ACROTECH

* ACROTECH BIOPHARMA LLC
 BELEODAQ, BELINOSTAT
 EVOMELA, MELPHALAN HYDROCHLORIDE
 FOLOTYN, PRALATREXATE
 FUSILEV, LEVOLEUCOVORIN CALCIUM
 KHAPZORY, LEVOLEUCOVORIN
 MARQIBO KIT, VINCRISTINE SULFATE

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 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
 DONEPEZIL HYDROCHLORIDE, DONEPEZIL 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
 ISOSORBIDE MONONITRATE, ISOSORBIDE MONONITRATE
 LAMOTRIGINE, LAMOTRIGINE
 LEVETIRACETAM, LEVETIRACETAM
 LOVASTATIN, LOVASTATIN
 METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
 METOPROLOL SUCCINATE, METOPROLOL SUCCINATE
 NIFEDIPINE, NIFEDIPINE
 OMEPRAZOLE AND SODIUM BICARBONATE, OMEPRAZOLE (OTC)
 OXAZEPAM, OXAZEPAM
 OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

- * ACTAVIS ELIZABETH LLC
 - OXYCODONE HYDROCHLORIDE AND IBUPROFEN, IBUPROFEN
 - OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
 - PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE
 - PROPRANOLOL HYDROCHLORIDE, PROPRANOLOL HYDROCHLORIDE
 - PROPYLTHIOURACIL, PROPYLTHIOURACIL
 - RANOLAZINE, RANOLAZINE
 - ROPINIROLE HYDROCHLORIDE, ROPINIROLE HYDROCHLORIDE
 - TEMAZEPAM, TEMAZEPAM
 - TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE
 - ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE
- * ACTAVIS ELIZABETH LLC AN INDIRECT WHOLLY OWNED SUB OF TEVA PHARMACEUTICALS USA INC
 - ALPRAZOLAM, ALPRAZOLAM
 - CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
 - LAMOTRIGINE, LAMOTRIGINE
 - MORPHINE SULFATE, MORPHINE SULFATE
 - OXYMORPHONE HYDROCHLORIDE, OXYMORPHONE HYDROCHLORIDE
 - PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE

ACTAVIS INC

- * ACTAVIS INC
 - DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE
 - DOXORUBICIN HYDROCHLORIDE, DOXORUBICIN HYDROCHLORIDE
 - METHOXSALLEN, METHOXSALLEN

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)
 - MESALAMINE, MESALAMINE
 - METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE

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
 - DESMOPRESSIN ACETATE, DESMOPRESSIN ACETATE
 - DICLOFENAC SODIUM AND MISOPROSTOL, DICLOFENAC SODIUM
 - DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
 - DIVALPROEX SODIUM, DIVALPROEX SODIUM
 - DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
 - DOXYLAMINE SUCCINATE AND PYRIDOXINE HYDROCHLORIDE, DOXYLAMINE SUCCINATE
 - DUTASTERIDE AND TAMSULOSIN HYDROCHLORIDE, DUTASTERIDE
 - HYDROCODONE BITARTRATE AND IBUPROFEN, HYDROCODONE BITARTRATE
 - LEVETIRACETAM, LEVETIRACETAM
 - LORATADINE, LORATADINE (OTC)
 - METAXALONE, METAXALONE
 - METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 - METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
 - METOPROLOL SUCCINATE, METOPROLOL SUCCINATE
 - NAPROXEN SODIUM, NAPROXEN SODIUM
 - NITROFURANTOIN, NITROFURANTOIN, MACROCRYSTALLINE
 - NITROGLYCERIN, NITROGLYCERIN
 - OMEPRAZOLE, OMEPRAZOLE
 - OXYCODONE AND ASPIRIN, ASPIRIN
 - PALIPERIDONE, PALIPERIDONE
 - PAROXETINE MESYLATE, PAROXETINE MESYLATE
 - POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
 - PREDNISONE, PREDNISONE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

**** A ****

- * ACTAVIS LABORATORIES FL INC
 - RAMELTEON, RAMELTEON
 - RISPERIDONE, RISPERIDONE
 - TAMOXIFEN CITRATE, TAMOXIFEN CITRATE
 - TAZTIA XT, DILTIAZEM HYDROCHLORIDE
 - TETRABENAZINE, TETRABENAZINE
 - TRANEXAMIC ACID, TRANEXAMIC ACID
 - TROSPIMUM CHLORIDE, TROSPIMUM CHLORIDE
 - VALGANCICLOVIR HYDROCHLORIDE, VALGANCICLOVIR HYDROCHLORIDE
 - ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE

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
 - FIORICET W/ CODEINE, ACETAMINOPHEN
 - LIDOCAINE, LIDOCAINE
 - PROGESTERONE, PROGESTERONE
 - TESTOSTERONE, TESTOSTERONE
- * ACTAVIS LABORATORIES UT INC INDIRECT WHOLLY OWNED SUB OF TEVA PHARMACEUTICALS USA INC
 - PIMECROLIMUS, PIMECROLIMUS
 - TESTOSTERONE, TESTOSTERONE

ACTAVIS LLC

- * ACTAVIS LLC
 - AZACITIDINE, AZACITIDINE
 - DAPSONE, DAPSONE
 - FLUDARABINE PHOSPHATE, FLUDARABINE PHOSPHATE
 - LEVOLEUCOVORIN CALCIUM, LEVOLEUCOVORIN CALCIUM
 - MELPHALAN HYDROCHLORIDE, MELPHALAN HYDROCHLORIDE
- * ACTAVIS LLC AN INDIRECT WHOLLY-OWNED SUB OF TEVA PHARMACEUTICALS USA INC
 - BUSULFAN, BUSULFAN
 - DOCETAXEL, DOCETAXEL
 - HYDROXOCOBALAMIN, HYDROXOCOBALAMIN
 - OXALIPLATIN, OXALIPLATIN

ACTAVIS MID ATLANTIC

- * ACTAVIS MID ATLANTIC LLC
 - ACYCLOVIR, ACYCLOVIR
 - ADAPALENE AND BENZOYL PEROXIDE, ADAPALENE
 - 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
 - ENULOSE, LACTULOSE
 - GRISEOFULVIN, GRISEOFULVIN, MICROSIZE
 - HYDROCODONE BITARTRATE AND HOMATROPINE METHYLBROMIDE, HOMATROPINE METHYLBROMIDE
 - HYDROCORTISONE BUTYRATE, HYDROCORTISONE BUTYRATE
 - HYDROCORTISONE, HYDROCORTISONE
 - LEVETIRACETAM, LEVETIRACETAM
 - NITROFURANTOIN, NITROFURANTOIN
 - NYSTATIN, NYSTATIN
 - PROMETH VC W/ CODEINE, CODEINE PHOSPHATE
 - PROMETHAZINE HYDROCHLORIDE AND CODEINE PHOSPHATE, CODEINE PHOSPHATE
 - 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

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

**** A ****

ACTAVIS TOTOWA

* ACTAVIS TOTOWA LLC
 DESIPRAMINE HYDROCHLORIDE, DESIPRAMINE HYDROCHLORIDE
 EPIRUBICIN HYDROCHLORIDE, EPIRUBICIN HYDROCHLORIDE
 FINASTERIDE, FINASTERIDE
 FLUDARABINE PHOSPHATE, FLUDARABINE PHOSPHATE
 IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE
 OXALIPLATIN, OXALIPLATIN
 PACLITAXEL, PACLITAXEL
 PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
 REPAGLINIDE, REPAGLINIDE
 TOPOTECAN HYDROCHLORIDE, TOPOTECAN HYDROCHLORIDE
 VINOURELBINE TARTRATE, VINOURELBINE TARTRATE

ACTAVIS TOTOWA TEVA

* ACTAVIS TOTOWA LLC AN INDIRECT WHOLLY OWNED SUB OF TEVA PHARMACEUTICALS USA INC
 FINASTERIDE, FINASTERIDE

ACTELION PHARMS

* ACTELION PHARMACEUTICALS LTD
 TRACLEER, BOSENTAN

ACTELION PHARMS LTD

* ACTELION PHARMACEUTICALS LTD
 OPSUMIT, MACITENTAN
 TRACLEER, BOSENTAN
 UPTRAVI, SELEXIPAG
 VELETRI, EPOPROSTENOL SODIUM
 VENTAVIS, ILOPROST
 ZAVESCA, MIGLUSTAT

ACTIENT PHARMS

* ACTIENT PHARMACEUTICALS LLC
 THEO-24, THEOPHYLLINE

ADAMAS PHARMA

* ADAMAS PHARMA LLC
 GOCOVRI, AMANTADINE HYDROCHLORIDE

ADAMIS PHARMS CORP

* ADAMIS PHARMACEUTICALS CORP
 SYMJEPI, EPINEPHRINE

ADAPT

* ADAPT PHARMA OPERATIONS LTD
 NARCAN, NALOXONE HYDROCHLORIDE

ADARE PHARMS INC

* ADARE PHARMACEUTICALS INC
 DEXMETHYLPHENIDATE HYDROCHLORIDE, DEXMETHYLPHENIDATE HYDROCHLORIDE
 POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
 PROPRANOLOL HYDROCHLORIDE, PROPRANOLOL HYDROCHLORIDE

ADDMEDICA SAS

* ADDMEDICA SAS
 SIKLOS, HYDROXYUREA

ADHERA

* ADHERA THERAPEUTICS INC
 PRESTALIA, AMLODIPINE BESYLATE

ADIENNE SA

* ADIENNE SA
 TEPADINA, THIOTEPA

AEGERION

* AEGERION PHARMACEUTICALS INC
 JUXTAPID, LOMITAPIDE MESYLATE

AERIE PHARMS INC

* AERIE PHARMACEUTICALS INC
 RHOPRESSA, NETARSUDIL MESYLATE
 ROCKLATAN, LATANOPROST

AGILE

* AGILE THERAPEUTICS INC
 TWIRLA, ETHINYL ESTRADIOL

AGIOS PHARMS INC

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

* AGIOS PHARMACEUTICALS INC
TIBSOVO, IVOSIDENIB

AGNITIO

* AGNITIO INC
ETHACRYNIC ACID, ETHACRYNIC ACID
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

AJANTA PHARMA LTD

* AJANTA PHARMA LTD
ALMOTRIPTAN MALATE, ALMOTRIPTAN MALATE
AMLODIPINE AND OLMESARTAN MEDOXOMIL, AMLODIPINE BESYLATE
ARIPIPIRAZOLE, ARIPIPIRAZOLE
CAPTOPRIL, CAPTOPRIL
CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
DIVALPROEX SODIUM, DIVALPROEX SODIUM
DOXEPIN HYDROCHLORIDE, DOXEPIN HYDROCHLORIDE
DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE
ELETRIPTAN HYDROBROMIDE, ELETRIPTAN HYDROBROMIDE
ENTACAPONE, ENTACAPONE
FENOFIBRATE (MICRONIZED), FENOFIBRATE
FENOFIBRATE, FENOFIBRATE
MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
MONTELUKAST SODIUM, MONTELUKAST SODIUM
OMEPRAZOLE AND SODIUM BICARBONATE, OMEPRAZOLE
OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE
RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
RANOLAZINE, RANOLAZINE
RISPERIDONE, RISPERIDONE
SILDENAFIL CITRATE, SILDENAFIL CITRATE
SILODOSIN, SILODOSIN
SOLIFENACIN SUCCINATE, SOLIFENACIN SUCCINATE
TADALAFIL, TADALAFIL
VALGANCICLOVIR HYDROCHLORIDE, VALGANCICLOVIR HYDROCHLORIDE
VORICONAZOLE, VORICONAZOLE
ZOLMITRIPTAN, ZOLMITRIPTAN

AKARX INC

* AKARX INC
DOPTELET, AVATROMBOPAG MALEATE

AKCEA THERAPY

* AKCEA THERAPEUTICS INC
TEGSEDI, INOTERSEN SODIUM

AKORN

* AKORN INC
ADENOSINE, ADENOSINE
AK-FLUOR 10%, FLUORESCEIN SODIUM
AK-FLUOR 25%, FLUORESCEIN SODIUM
AKBETA, LEVOBUNOLOL HYDROCHLORIDE
AKPENTOLATE, CYCLOPENTOLATE HYDROCHLORIDE
AKTEN, LIDOCAINE HYDROCHLORIDE
AKTOB, TOBRAMYCIN
ALFENTA, ALFENTANIL HYDROCHLORIDE
ATROPINE SULFATE, ATROPINE SULFATE
AZELASTINE HYDROCHLORIDE, AZELASTINE HYDROCHLORIDE
BACITRACIN ZINC AND POLYMYXIN B SULFATE, BACITRACIN ZINC
BACITRACIN, BACITRACIN
BAL, DIMERCAPROL
BALANCED SALT, CALCIUM CHLORIDE
BETAXOLOL HYDROCHLORIDE, BETAXOLOL HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ******* AKORN INC**

BRIMONIDINE TARTRATE, BRIMONIDINE TARTRATE
 CALCITRIOL, CALCITRIOL
 CAPASTAT SULFATE, CAPREOMYCIN SULFATE
 CARBOPLATIN, CARBOPLATIN
 CICLOPIROX, CICLOPIROX
 CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
 CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
 CROMOLYN SODIUM, CROMOLYN SODIUM
 DEMECLOCYCLINE HYDROCHLORIDE, DEMECLOCYCLINE HYDROCHLORIDE
 DESOXIMETASONE, DESOXIMETASONE
 DICLOFENAC SODIUM, DICLOFENAC SODIUM
 DIFLORASONE DIACETATE, DIFLORASONE DIACETATE
 EPINASTINE HYDROCHLORIDE, EPINASTINE HYDROCHLORIDE
 EPTIFIBATIDE, EPTIFIBATIDE
 ERYTHROMYCIN, ERYTHROMYCIN
 ETHAMBUTOL HYDROCHLORIDE, ETHAMBUTOL HYDROCHLORIDE
 ETHOSUXIMIDE, ETHOSUXIMIDE
 FLUOCINOLONE ACETONIDE, FLUOCINOLONE ACETONIDE
 GENTAK, GENTAMICIN SULFATE
 GENTAMICIN SULFATE, GENTAMICIN SULFATE
 HALOPERIDOL, HALOPERIDOL LACTATE
 HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
 HYDROMORPHONE HYDROCHLORIDE, HYDROMORPHONE HYDROCHLORIDE
 IC-GREEN, INDOCYANINE GREEN
 IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE
 KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
 KETOTIFEN FUMARATE, KETOTIFEN FUMARATE (OTC)
 LATANOPROST, LATANOPROST
 LEVOFLOXACIN, LEVOFLOXACIN
 LIDOCAINE HYDROCHLORIDE, LIDOCAINE HYDROCHLORIDE
 LORAZEPAM, LORAZEPAM
 METHADONE HYDROCHLORIDE, METHADONE HYDROCHLORIDE
 MOXIFLOXACIN HYDROCHLORIDE, MOXIFLOXACIN HYDROCHLORIDE
 NALOXONE HYDROCHLORIDE, NALOXONE HYDROCHLORIDE
 NEDOCROMIL SODIUM, NEDOCROMIL SODIUM
 NEOMYCIN AND POLYMYXIN B SULFATES AND BACITRACIN ZINC, BACITRACIN ZINC
 NEOMYCIN AND POLYMYXIN B SULFATES, BACITRACIN ZINC AND HYDROCORTISONE, BACITRACIN ZINC
 NYSTATIN AND TRIAMCINOLONE ACETONIDE, NYSTATIN
 OFLOXACIN, OFLOXACIN
 OLOPATADINE HYDROCHLORIDE, OLOPATADINE HYDROCHLORIDE
 ORPHENADRINE CITRATE, ORPHENADRINE CITRATE
 PAREMYD, HYDROXYAMPHETAMINE HYDROBROMIDE
 PARICALCITOL, PARICALCITOL
 PYRAZINAMIDE, PYRAZINAMIDE
 RIFAMPIN, RIFAMPIN
 ROPIVACAINE HYDROCHLORIDE, ROPIVACAINE HYDROCHLORIDE
 SODIUM NITROPRUSSIDE, SODIUM NITROPRUSSIDE
 SUBLIMAZE PRESERVATIVE FREE, FENTANYL CITRATE
 SUFENTA PRESERVATIVE FREE, SUFENTANIL CITRATE
 SULFACETAMIDE SODIUM, SULFACETAMIDE SODIUM
 TERBUTALINE SULFATE, TERBUTALINE SULFATE
 TIMOLOL MALEATE, TIMOLOL MALEATE
 TIMOLOL, TIMOLOL
 TOBRAMYCIN SULFATE, TOBRAMYCIN SULFATE
 TRANEXAMIC ACID, TRANEXAMIC ACID
 TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE
 TROPICACYL, TROPICAMIDE
 VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE
 VORICONAZOLE, VORICONAZOLE
 ZOLEDRONIC ACID, ZOLEDRONIC ACID

AKORN INC*** AKORN INC**

ACETYLCYSTEINE, ACETYLCYSTEINE
 APRACLONIDINE HYDROCHLORIDE, APRACLONIDINE HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ******* AKORN INC**

CEFTRIAZONE, CEFTRIAZONE SODIUM
 CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE
 CLINDAMYCIN PHOSPHATE IN 5% DEXTROSE IN PLASTIC CONTAINER, CLINDAMYCIN PHOSPHATE
 CYCLOPENTOLATE HYDROCHLORIDE, CYCLOPENTOLATE HYDROCHLORIDE
 DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE
 DICYCLOMINE HYDROCHLORIDE, DICYCLOMINE HYDROCHLORIDE
 DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
 DORZOLAMIDE HYDROCHLORIDE AND TIMOLOL MALEATE, DORZOLAMIDE HYDROCHLORIDE
 DOXERCALCIFEROL, DOXERCALCIFEROL
 DRONABINOL, DRONABINOL
 EPHEDRINE SULFATE, EPHEDRINE SULFATE
 EPIRUBICIN HYDROCHLORIDE, EPIRUBICIN HYDROCHLORIDE
 GRANISETRON HYDROCHLORIDE, GRANISETRON HYDROCHLORIDE
 INAPSINE, DROPERIDOL
 LABETALOL HYDROCHLORIDE, LABETALOL HYDROCHLORIDE
 MIDAZOLAM HYDROCHLORIDE, MIDAZOLAM HYDROCHLORIDE
 MYCOPHENOLATE MOFETIL HYDROCHLORIDE, MYCOPHENOLATE MOFETIL HYDROCHLORIDE
 NAPHAZOLINE HYDROCHLORIDE AND PHENIRAMINE MALEATE, NAPHAZOLINE HYDROCHLORIDE (OTC)
 OLOPATADINE HYDROCHLORIDE, OLOPATADINE HYDROCHLORIDE
 PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
 PHENYLEPHRINE HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE
 PILOCARPINE HYDROCHLORIDE, PILOCARPINE HYDROCHLORIDE
 PROPARACAINE HYDROCHLORIDE, PROPARACAINE HYDROCHLORIDE
 ROPIVACAINE HYDROCHLORIDE, ROPIVACAINE HYDROCHLORIDE
 TOBRAMYCIN, TOBRAMYCIN
 TRIMETHOPRIM SULFATE AND POLYMYXIN B SULFATE, POLYMYXIN B SULFATE
 ZOLEDRONIC ACID, ZOLEDRONIC ACID

ALAN LABS INC

* ALAN LABORATORIES INC
 DARIFENACIN HYDROBROMIDE, DARIFENACIN HYDROBROMIDE

ALCON

* ALCON LABORATORIES INC
 BSS PLUS, CALCIUM CHLORIDE
 BSS, CALCIUM CHLORIDE
 MIostat, CARBACHOL
 NAPHCN-A, NAPHAZOLINE HYDROCHLORIDE (OTC)

ALCON LABS

* ALCON LABORATORIES LTD
 TETRACAINE HYDROCHLORIDE, TETRACAINE HYDROCHLORIDE

ALCON LABS INC

* ALCON LABORATORIES INC
 ALCAINE, PROPARACAINE HYDROCHLORIDE
 CYCLOGLY, CYCLOPENTOLATE HYDROCHLORIDE
 CYCLOMYDRIL, CYCLOPENTOLATE HYDROCHLORIDE
 FLUORESCITE, FLUORESCIEIN SODIUM
 ISOPTO ATROPINE, ATROPINE SULFATE

ALCON PHARMS LTD

* ALCON PHARMACEUTICALS LTD
 BETADINE, POVIDONE-IODINE
 ZADITOR, KETOTIFEN FUMARATE (OTC)

ALEMBIC LTD

* ALEMBIC LTD
 LITHIUM CARBONATE, LITHIUM CARBONATE
 ROPINIROLE HYDROCHLORIDE, ROPINIROLE HYDROCHLORIDE

ALEMBIC PHARMS LTD

* ALEMBIC PHARMACEUTICALS LTD
 ACETAZOLAMIDE, ACETAZOLAMIDE
 ACYCLOVIR, ACYCLOVIR
 AMANTADINE HYDROCHLORIDE, AMANTADINE HYDROCHLORIDE
 AMLODIPINE AND OLMESARTAN MEDOXOMIL, AMLODIPINE BESYLATE
 AMLODIPINE BESYLATE AND VALSARTAN, AMLODIPINE BESYLATE
 ARIPIPRAZOLE, ARIPIPRAZOLE
 AZELASTINE HYDROCHLORIDE, AZELASTINE HYDROCHLORIDE
 AZITHROMYCIN, AZITHROMYCIN

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

* ALEMBIC PHARMACEUTICALS LTD
 BIMATOPROST, BIMATOPROST
 BOSENTAN, BOSENTAN
 BROMFENAC SODIUM, BROMFENAC SODIUM
 BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
 CANDESARTAN CILEXETIL, CANDESARTAN CILEXETIL
 CARBIDOPA AND LEVODOPA, CARBIDOPA
 CELECOXIB, CELECOXIB
 CLONAZEPAM, CLONAZEPAM
 CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
 DARIFENACIN HYDROBROMIDE, DARIFENACIN HYDROBROMIDE
 DEFERASIROX, DEFERASIROX
 DESVENLAFAXINE SUCCINATE, DESVENLAFAXINE SUCCINATE
 DESVENLAFAXINE, DESVENLAFAXINE
 DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
 DORZOLAMIDE HYDROCHLORIDE, DORZOLAMIDE HYDROCHLORIDE
 DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
 DOXYCYCLINE, DOXYCYCLINE
 DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE
 FAMOTIDINE, FAMOTIDINE
 FEBUXOSTAT, FEBUXOSTAT
 FENOFIBRATE, FENOFIBRATE
 FENOFIBRIC ACID, CHOLINE FENOFIBRATE
 FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 IRBESARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 IRBESARTAN, IRBESARTAN
 ITRACONAZOLE, ITRACONAZOLE
 LAMOTRIGINE, LAMOTRIGINE
 LEFLUNOMIDE, LEFLUNOMIDE
 LINEZOLID, LINEZOLID
 LITHIUM CARBONATE, LITHIUM CARBONATE
 LOSARTAN POTASSIUM AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 LOSARTAN POTASSIUM, LOSARTAN POTASSIUM
 MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
 MEPROBAMATE, MEPROBAMATE
 METOPROLOL TARTRATE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 METOPROLOL TARTRATE, METOPROLOL TARTRATE
 METRONIDAZOLE, METRONIDAZOLE
 MODAFINIL, MODAFINIL
 MOXIFLOXACIN HYDROCHLORIDE, MOXIFLOXACIN HYDROCHLORIDE
 OLMESARTAN MEDOXOMIL AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 OLMESARTAN MEDOXOMIL, OLMESARTAN MEDOXOMIL
 OLOPATADINE HYDROCHLORIDE, OLOPATADINE HYDROCHLORIDE
 OSELTAMIVIR PHOSPHATE, OSELTAMIVIR PHOSPHATE
 PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE
 PREGABALIN, PREGABALIN
 QUETIAPINE FUMARATE, QUETIAPINE FUMARATE
 RIVASTIGMINE TARTRATE, RIVASTIGMINE TARTRATE
 ROPINIROLE HYDROCHLORIDE, ROPINIROLE HYDROCHLORIDE
 SILODOSIN, SILODOSIN
 SOLIFENACIN SUCCINATE, SOLIFENACIN SUCCINATE
 TADALAFIL, TADALAFIL
 TELMISARTAN AND AMLODIPINE, AMLODIPINE BESYLATE
 TELMISARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 TELMISARTAN, TELMISARTAN
 TEMAZEPAM, TEMAZEPAM
 TERIFLUNOMIDE, TERIFLUNOMIDE
 THEOPHYLLINE, THEOPHYLLINE
 TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE
 TOBRAMYCIN, TOBRAMYCIN
 TRAVOPROST, TRAVOPROST
 VALSARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 VALSARTAN, VALSARTAN
 VARDENAFIL HYDROCHLORIDE, VARDENAFIL HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

**** A ****

* ALEMBIC PHARMACEUTICALS LTD
 VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE
 VILAZODONE HYDROCHLORIDE, VILAZODONE HYDROCHLORIDE
 ZOLMITRIPTAN, ZOLMITRIPTAN

ALEOR DERMACEUTICALS

* ALEOR DERMACEUTICALS LTD
 CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
 DESONIDE, DESONIDE
 LIDOCAINE, LIDOCAINE

ALEXZA PHARMS

* ALEXZA PHARMACEUTICALS INC
 ADASUVE, LOXAPINE

ALFASIGMA

* ALFASIGMA USA INC
 ZELNORM, TEGASEROD MALEATE

ALIGNSCIENCE PHARMA

* ALIGNSCIENCE PHARMA INC
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 QUETIAPINE FUMARATE, QUETIAPINE FUMARATE

ALIMERA SCIENCES INC

* ALIMERA SCIENCES INC
 ILUVIEN, 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
 ARIPIPRAZOLE, ARIPIPRAZOLE
 AZELASTINE HYDROCHLORIDE, AZELASTINE HYDROCHLORIDE
 CAPECITABINE, CAPECITABINE
 CEFIXIME, CEFIXIME
 CEFUROXIME AXETIL, CEFUROXIME AXETIL
 CEPHALEXIN, CEPHALEXIN
 CHLORTHALIDONE, CHLORTHALIDONE
 CINACALCET HYDROCHLORIDE, CINACALCET HYDROCHLORIDE
 CLOBAZAM, CLOBAZAM
 COLCHICINE, COLCHICINE
 COLESEVELAM HYDROCHLORIDE, COLESEVELAM HYDROCHLORIDE
 DABIGATRAN ETEXILATE MESYLATE, DABIGATRAN ETEXILATE MESYLATE
 DALFAMPRIDINE, DALFAMPRIDINE
 DEFERASIROX, DEFERASIROX
 DEXMETHYLPHENIDATE HYDROCHLORIDE, DEXMETHYLPHENIDATE HYDROCHLORIDE
 DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE
 ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM
 EZETIMIBE AND SIMVASTATIN, EZETIMIBE
 EZETIMIBE, EZETIMIBE
 FINASTERIDE, FINASTERIDE
 GABAPENTIN, GABAPENTIN
 HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
 ITRACONAZOLE, ITRACONAZOLE
 LAMOTRIGINE, LAMOTRIGINE
 LANSOPRAZOLE, LANSOPRAZOLE
 LIDOCAINE, LIDOCAINE
 LINEZOLID, LINEZOLID
 MARINOL, DRONABINOL
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
 MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE
 MYCOPHENOLATE MOFETIL, MYCOPHENOLATE MOFETIL

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

* ALKEM LABORATORIES LTD
 OLANZAPINE, OLANZAPINE
 OLMESARTAN MEDOXOMIL, OLMESARTAN MEDOXOMIL
 OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
 PREGABALIN, PREGABALIN
 QUETIAPINE FUMARATE, QUETIAPINE FUMARATE
 RABEPRAZOLE SODIUM, RABEPRAZOLE SODIUM
 RASAGILINE MESYLATE, RASAGILINE MESYLATE
 RILUZOLE, RILUZOLE
 RIZATRIPTAN BENZOATE, RIZATRIPTAN BENZOATE
 ROSUVASTATIN CALCIUM, ROSUVASTATIN CALCIUM
 SILDENAFIL CITRATE, SILDENAFIL CITRATE
 SOLIFENACIN SUCCINATE, SOLIFENACIN SUCCINATE
 TAMSULOSIN HYDROCHLORIDE, TAMSULOSIN HYDROCHLORIDE
 TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE
 TRAMADOL HYDROCHLORIDE AND ACETAMINOPHEN, ACETAMINOPHEN
 TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE
 VALSARTAN, VALSARTAN

ALKERMES

* ALKERMES INC
 VIVITROL, NALTREXONE

ALKERMES INC

* ALKERMES INC
 ARISTADA INITIO KIT, ARIPIPRAZOLE LAUROXIL
 ARISTADA, ARIPIPRAZOLE LAUROXIL

ALLEGIANCE HLTHCARE

* ALLEGIANCE HEALTHCARE CORP
 POVIDONE IODINE, POVIDONE-IODINE (OTC)

ALLEGIS

* ALLEGIS HOLDINGS LLC
 PRIMOSOL, TRIMETHOPRIM HYDROCHLORIDE

ALLERGAN

* ALLERGAN
 ACULAR LS, KETOROLAC TROMETHAMINE
 ALPHAGAN P, BRIMONIDINE TARTRATE
 BLEPH-10, SULFACETAMIDE SODIUM
 GENOPTIC, GENTAMICIN SULFATE
 ZYMAXID, GATIFLOXACIN

* ALLERGAN INC
 ACULAR, KETOROLAC TROMETHAMINE
 ACUVAIL, KETOROLAC TROMETHAMINE
 ACZONE, DAPSONE
 ALOCRIL, NEDOCROMIL SODIUM
 ALPHAGAN P, BRIMONIDINE TARTRATE
 AVAGE, TAZAROTENE
 COMBIGAN, BRIMONIDINE TARTRATE
 ELESTAT, EPINASTINE HYDROCHLORIDE
 LASTACFT, ALCAFTADINE
 LATISSE, BIMATOPROST
 LUMIGAN, BIMATOPROST
 OCUFLOX, OFLOXACIN
 OZURDEX, DEXAMETHASONE
 POLYTRIM, POLYMYXIN B SULFATE
 RESTASIS MULTIDOSE, CYCLOSPORINE
 RESTASIS, CYCLOSPORINE
 TAZORAC, TAZAROTENE
 ZYMAR, GATIFLOXACIN

* ALLERGAN PHARMACEUTICAL
 BETAGAN, LEVOBUNOLOL HYDROCHLORIDE
 BLEPHAMIDE S.O.P., PREDNISOLONE ACETATE
 BLEPHAMIDE, PREDNISOLONE ACETATE
 FML FORTE, FLUOROMETHOLONE
 FML, FLUOROMETHOLONE
 OCUFEN, FLURBIPROFEN SODIUM
 PRED FORTE, PREDNISOLONE ACETATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

* ALLERGAN PHARMACEUTICAL
 PRED MILD, PREDNISOLONE ACETATE
 PRED-G, GENTAMICIN SULFATE

* ALLERGAN SALES LLC
 ACTIGALL, URSODIOL
 ALORA, ESTRADIOL
 ANDRODERM, TESTOSTERONE
 AVYCAZ, AVIBACTAM SODIUM
 BENTYL PRESERVATIVE FREE, DICYCLOMINE HYDROCHLORIDE
 BENTYL, DICYCLOMINE HYDROCHLORIDE
 BREVICON 28-DAY, ETHINYL ESTRADIOL
 BYSTOLIC, NEBIVOLOL HYDROCHLORIDE
 CANASA, MESALAMINE
 CARAFATE, SUCRALFATE
 CELEXA, CITALOPRAM HYDROBROMIDE
 CONDYLOX, PODOFILOX
 CRINONE, PROGESTERONE
 DALVANCE, DALBAVANCIN HYDROCHLORIDE
 ESTRACE, ESTRADIOL
 FETZIMA, LEVOMILNACIPRAN HYDROCHLORIDE
 FIORINAL W/CODEINE, ASPIRIN
 FIORINAL, ASPIRIN
 GELNIQUE, OXYBUTYNIN CHLORIDE
 INFED, IRON DEXTRAN
 KADIAN, MORPHINE SULFATE
 LEXAPRO, ESCITALOPRAM OXALATE
 LINZESS, LINACLOTIDE
 MICROZIDE, HYDROCHLOROTHIAZIDE
 NAMENDA, MEMANTINE HYDROCHLORIDE
 NAMZARIC, DONEPEZIL HYDROCHLORIDE
 NORINYL 1+35 21-DAY, ETHINYL ESTRADIOL
 NORINYL 1+35 28-DAY, ETHINYL ESTRADIOL
 OXYTROL FOR WOMEN, OXYBUTYNIN (OTC)
 OXYTROL, OXYBUTYNIN
 PYLERA, BISMUTH SUBCITRATE POTASSIUM
 RAPAFLO, SILODOSIN
 RECTIV, NITROGLYCERIN
 SAPHRIS, ASENAPINE MALEATE
 SAVELLA, MILNACIPRAN HYDROCHLORIDE
 TEFLARO, CEFTAROLINE FOSAMIL
 TRELSTAR, TRIPTORELIN PAMOATE
 UBRELVY, UBROGEPANT
 URSO 250, URSODIOL
 URSO FORTE, URSODIOL
 VIIBRYD, VILAZODONE HYDROCHLORIDE
 VRAYLAR, CARIPRAZINE HYDROCHLORIDE

ALLERGAN HOLDINGS

* ALLERGAN HOLDINGS UNLTD CO
 VIBERZI, ELUXADOLINE

ALLERQUEST

* ALLERQUEST LLC
 PRE-PEN, BENZYL PENICILLOYL POLYLYSINE

ALLIED

* ALLIED PHARMA INC
 CARISOPRODOL, CARISOPRODOL
 DROSPIRENONE AND ETHINYL ESTRADIOL, DROSPIRENONE
 ROSUVASTATIN CALCIUM, ROSUVASTATIN CALCIUM
 TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE

ALMATICA

* ALMATICA PHARMA INC
 GRALISE, GABAPENTIN

ALMIRALL

* ALMIRALL LLC
 ACTICLATE, DOXYCYCLINE HYCLATE
 ACZONE, DAPSONE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

**** A ****

* ALMIRALL LLC

ALTABAX, RETAPAMULIN
 AZELEX, AZELAIC ACID
 CORDRAN SP, FLURANDRENOLIDE
 CORDRAN, FLURANDRENOLIDE
 FLUOROPLEX, FLUOROURACIL
 SEYSARA, SARECYCLINE HYDROCHLORIDE
 VELTIN, CLINDAMYCIN PHOSPHATE
 VERDESO, DESONIDE
 XOLEGEL, KETOCONAZOLE

ALNYLAM PHARMS INC

* ALNYLAM PHARMACEUTICALS INC
 GIVLAARI, GIVOSIRAN SODIUM
 ONPATTRO, PATISIRAN SODIUM

ALRA

* ALRA LABORATORIES INC
 CHOLAC, LACTULOSE
 CONSTILAC, LACTULOSE
 GEN-XENE, CLORAZEPATE DIPOTASSIUM
 IBU-TAB 200, IBUPROFEN (OTC)
 IBU-TAB, IBUPROFEN

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

ALVOGEN

* ALVOGEN GROUP HOLDINGS 2 LLC
 DAPSONE, DAPSONE
 ETHACRYNIC ACID, ETHACRYNIC ACID
 OFLOXACIN, OFLOXACIN

* ALVOGEN GROUP HOLDINGS 3 LLC
 FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 FORFIVO XL, BUPROPION HYDROCHLORIDE

* ALVOGEN GROUP HOLDINGS 4 LLC
 THYRO-TABS, LEVOTHYROXINE SODIUM **

* ALVOGEN GROUP HOLDINGS LLC
 ADALAT CC, NIFEDIPINE

* ALVOGEN INC
 TRAZODONE HYDROCHLORIDE, TRAZODONE HYDROCHLORIDE

* ALVOGEN MALTA OPERATIONS LTD
 ATENOLOL AND CHLORTHALIDONE, ATENOLOL
 ATENOLOL, ATENOLOL
 BONSIITY, TERIPARATIDE
 BUDESONIDE, BUDESONIDE
 CARBIDOPA, CARBIDOPA
 DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE
 DISULFIRAM, DISULFIRAM
 EXEMESTANE, EXEMESTANE
 FELBAMATE, FELBAMATE
 HYDROCODONE BITARTRATE, HYDROCODONE BITARTRATE
 MACROBID, NITROFURANTOIN
 MACRODANTIN, NITROFURANTOIN, MACROCRYSTALLINE
 MELPHALAN, MELPHALAN
 NAPRELAN, NAPROXEN SODIUM
 NATEGLINIDE, NATEGLINIDE
 NEVIRAPINE, NEVIRAPINE
 NITROGLYCERIN, NITROGLYCERIN
 OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN
 OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

* ALVOGEN MALTA OPERATIONS LTD
 PYRIDOSTIGMINE BROMIDE, PYRIDOSTIGMINE BROMIDE
 RIVASTIGMINE, RIVASTIGMINE
 SPECTAZOLE, ECONAZOLE NITRATE
 TENORETIC 100, ATENOLOL
 TENORETIC 50, ATENOLOL
 TENORMIN, ATENOLOL
 VORICONAZOLE, VORICONAZOLE
 ZESTORETIC, HYDROCHLOROTHIAZIDE
 ZESTRIL, LISINOPRIL

ALVOGEN INC

* ALVOGEN INC
 ACETYLCYSTEINE, ACETYLCYSTEINE
 CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
 MELPHALAN HYDROCHLORIDE, MELPHALAN HYDROCHLORIDE
 PRIMAQUINE PHOSPHATE, PRIMAQUINE PHOSPHATE

ALVOGEN PINE BROOK

* ALVOGEN PINE BROOK LLC
 BUPRENORPHINE HYDROCHLORIDE AND NALOXONE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE
 ESTRADIOL, ESTRADIOL
 METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
 OSELTAMIVIR PHOSPHATE, OSELTAMIVIR PHOSPHATE

AM ANTIBIOTICS

* AMERICAN ANTIBIOTICS INC
 AMOXICILLIN, AMOXICILLIN

AM REGENT

* AMERICAN REGENT INC
 ACETYLCYSTEINE, ACETYLCYSTEINE
 BETAMETHASONE ACETATE AND BETAMETHASONE SODIUM PHOSPHATE, BETAMETHASONE ACETATE
 BUPRENORPHINE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE
 CAFFEINE CITRATE, CAFFEINE CITRATE
 CALCIUM CHLORIDE 10%, CALCIUM CHLORIDE
 CHLOROTHIAZIDE SODIUM, CHLOROTHIAZIDE SODIUM
 CYANOCOBALAMIN, CYANOCOBALAMIN
 CYCLOSPORINE, CYCLOSPORINE
 DESMOPRESSIN ACETATE, DESMOPRESSIN ACETATE
 DICYCLOMINE HYDROCHLORIDE, DICYCLOMINE HYDROCHLORIDE
 DROPERIDOL, DROPERIDOL
 ESTRADIOL VALERATE, ESTRADIOL VALERATE
 FOMEPIZOLE, FOMEPIZOLE
 GLYCOPYRROLATE, GLYCOPYRROLATE
 HYDROXYPROGESTERONE CAPROATE, HYDROXYPROGESTERONE CAPROATE
 HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE
 INJECTAFER, FERRIC CARBOXYMALTOSIDE
 LEVOCARNITINE, LEVOCARNITINE
 METHOCARBAMOL, METHOCARBAMOL
 METHYLERGONOVINE MALEATE, METHYLERGONOVINE MALEATE
 NEOSTIGMINE METHYLSULFATE, NEOSTIGMINE METHYLSULFATE
 NITROGLYCERIN, NITROGLYCERIN
 OLANZAPINE, OLANZAPINE
 SELENIOUS ACID, SELENIOUS ACID
 TESTOSTERONE CYPIONATE, TESTOSTERONE CYPIONATE
 TRANEXAMIC ACID, TRANEXAMIC ACID
 VENOFER, IRON SUCROSE
 ZINC SULFATE, ZINC SULFATE

AMAG PHARMA USA

* AMAG PHARMA USA INC
 MAKENA (AUTOINJECTOR), HYDROXYPROGESTERONE CAPROATE
 MAKENA PRESERVATIVE FREE, HYDROXYPROGESTERONE CAPROATE
 MAKENA, HYDROXYPROGESTERONE CAPROATE

AMAG PHARMS INC

* AMAG PHARMACEUTICALS INC
 FERAHEME, FERUMOXYTOL
 INTRAROSA, PRASTERONE
 VYLEESI (AUTOINJECTOR), BREMELANOTIDE ACETATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ******AMARIN PHARMS**

* AMARIN PHARMACEUTICALS IRELAND LTD
VASCEPA, ICOSAPENT ETHYL

AMERIGEN PHARMS LTD

* AMERIGEN PHARMACEUTICALS LTD
MIGLUSTAT, MIGLUSTAT
TEMOZOLOMIDE, TEMOZOLOMIDE

AMGEN

* AMGEN INC
SENSIPAR, CINACALCET HYDROCHLORIDE

AMGEN INC

* AMGEN INC
CORLANOR, IVABRADINE
CORLANOR, IVABRADINE HYDROCHLORIDE
OTEZLA, APREMILAST

AMICI

* AMICI PHARMACEUTICALS LLC
DICLOFENAC POTASSIUM, DICLOFENAC POTASSIUM

AMICUS THERAPS US

* AMICUS THERAPEUTICS US INC
GALAFOLD, MIGALASTAT HYDROCHLORIDE

AMNEAL

* AMNEAL EU LTD
AMINOCAPROIC ACID, AMINOCAPROIC ACID
BUSULFAN, BUSULFAN
CARMUSTINE, CARMUSTINE
CLOFARABINE, CLOFARABINE
CYCLOPHOSPHAMIDE, CYCLOPHOSPHAMIDE
DEFERASIROX, DEFERASIROX
DEXAMETHASONE SODIUM PHOSPHATE PRESERVATIVE FREE, DEXAMETHASONE SODIUM PHOSPHATE
DEXAMETHASONE SODIUM PHOSPHATE, DEXAMETHASONE SODIUM PHOSPHATE
DOCETAXEL, DOCETAXEL
DOXERCALCIFEROL, DOXERCALCIFEROL
DOXORUBICIN HYDROCHLORIDE, DOXORUBICIN HYDROCHLORIDE
EPHEDRINE SULFATE, EPHEDRINE SULFATE
FOSPHENYTOIN SODIUM, FOSPHENYTOIN SODIUM
FULVESTRANT, FULVESTRANT
GLYCOPYRROLATE, GLYCOPYRROLATE
ISOPROTERENOL HYDROCHLORIDE, ISOPROTERENOL HYDROCHLORIDE
LEVOLEUCOVORIN CALCIUM, LEVOLEUCOVORIN CALCIUM
METHYLPREDNISOLONE SODIUM SUCCINATE, METHYLPREDNISOLONE SODIUM SUCCINATE
NEOSTIGMINE METHYLSULFATE, NEOSTIGMINE METHYLSULFATE
NOREPINEPHRINE BITARTRATE, NOREPINEPHRINE BITARTRATE
OFLOXACIN, OFLOXACIN
PHENYLEPHRINE HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE
SODIUM NITROPRUSSIDE, SODIUM NITROPRUSSIDE
SUCCINYLCHOLINE CHLORIDE, SUCCINYLCHOLINE CHLORIDE
TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE
VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE

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

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

- * AMNEAL PHARMACEUTICALS
 - CALCITRIOL, CALCITRIOL
 - CALCIUM ACETATE, CALCIUM ACETATE
 - CAPECITABINE, CAPECITABINE
 - 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
 - DIVALPROEX SODIUM, DIVALPROEX SODIUM
 - ENTECAVIR, ENTECAVIR
 - ESCITALOPRAM OXALATE, ESCITALOPRAM OXALATE
 - ESTRADIOL, ESTRADIOL
 - FELBAMATE, FELBAMATE
 - GABAPENTIN, GABAPENTIN
 - HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
 - INDOMETHACIN, INDOMETHACIN
 - ITRACONAZOLE, ITRACONAZOLE
 - LEVETIRACETAM, LEVETIRACETAM
 - LIDOCAINE, LIDOCAINE
 - LINEZOLID, LINEZOLID
 - LORAZEPAM, LORAZEPAM
 - MEROPENEM, MEROPENEM
 - METAXALONE, METAXALONE
 - MOMETASONE FUROATE, MOMETASONE FUROATE
 - MONTELUKAST SODIUM, MONTELUKAST SODIUM
 - NIACIN, NIACIN
 - NITROFURANTOIN (MONOHYDRATE/MACROCRYSTALS), NITROFURANTOIN
 - NITROFURANTOIN, NITROFURANTOIN
 - NIZATIDINE, NIZATIDINE
 - NORETHINDRONE ACETATE, NORETHINDRONE ACETATE
 - ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
 - OXCARBAZEPINE, OXCARBAZEPINE
 - OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE
 - OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
 - PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
 - POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
 - PROMETHAZINE HYDROCHLORIDE AND CODEINE PHOSPHATE, CODEINE PHOSPHATE
 - PROMETHAZINE HYDROCHLORIDE AND PHENYLEPHRINE HYDROCHLORIDE, PHENYLEPHRINE
 - PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
 - PROMETHAZINE HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE W/CODEINE PHOSPHATE, CODEINE
 - QUININE SULFATE, QUININE SULFATE
 - RABEPRAZOLE SODIUM, RABEPRAZOLE SODIUM
 - RALOXIFENE HYDROCHLORIDE, RALOXIFENE HYDROCHLORIDE
 - RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
 - RISPERIDONE, RISPERIDONE
 - SILDENAFIL CITRATE, SILDENAFIL CITRATE
 - TELMISARTAN, TELMISARTAN
 - TEMAZEPAM, TEMAZEPAM
 - TEMOZOLOMIDE, TEMOZOLOMIDE
 - TRAMADOL HYDROCHLORIDE AND ACETAMINOPHEN, ACETAMINOPHEN
 - VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE
 - VORICONAZOLE, VORICONAZOLE
 - 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

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

* AMNEAL PHARMACEUTICALS OF NEW YORK LLC
 BUDESONIDE, BUDESONIDE
 BUPRENORPHINE HYDROCHLORIDE AND NALOXONE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE
 CELECOXIB, CELECOXIB
 DUTASTERIDE, DUTASTERIDE
 EPTIFIBATIDE, EPTIFIBATIDE
 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
 METHOTREXATE SODIUM, METHOTREXATE SODIUM
 METHYLERGONOVINE MALEATE, METHYLERGONOVINE MALEATE
 METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
 NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL AND FERROUS FUMARATE, ETHINYL ESTRADIOL
 NORETHINDRONE AND ETHINYL ESTRADIOL AND FERROUS FUMARATE, ETHINYL ESTRADIOL
 NORETHINDRONE, NORETHINDRONE
 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
 TOBRAMYCIN, TOBRAMYCIN
 TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE
 VALSARTAN, VALSARTAN
 VIGABATRIN, VIGABATRIN

AMNEAL PHARMS CO

* AMNEAL PHARMACEUTICALS CO GMBH
 ALBUTEROL SULFATE, ALBUTEROL SULFATE
 ARGATROBAN, ARGATROBAN
 BOSENTAN, BOSENTAN
 BUMETANIDE, BUMETANIDE
 BUSPIRONE HYDROCHLORIDE, BUSPIRONE HYDROCHLORIDE
 CHLORPROMAZINE HYDROCHLORIDE, CHLORPROMAZINE HYDROCHLORIDE
 CHLORZALIDONE, CHLORZALIDONE
 CLOBAZAM, CLOBAZAM
 CLOMIPRAMINE HYDROCHLORIDE, CLOMIPRAMINE HYDROCHLORIDE
 DESIPRAMINE HYDROCHLORIDE, DESIPRAMINE HYDROCHLORIDE
 DOXEPIN HYDROCHLORIDE, DOXEPIN HYDROCHLORIDE
 DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
 ELETRIPTAN HYDROBROMIDE, ELETRIPTAN HYDROBROMIDE
 ERYTHROMYCIN, ERYTHROMYCIN
 ETHACRYNIC ACID, ETHACRYNIC ACID
 ETODOLAC, ETODOLAC
 EZETIMIBE AND SIMVASTATIN, EZETIMIBE
 EZETIMIBE, EZETIMIBE
 FROVATRIPTAN SUCCINATE, FROVATRIPTAN SUCCINATE
 FUROSEMIDE, FUROSEMIDE
 HYDROXYCHLOROQUINE SULFATE, HYDROXYCHLOROQUINE SULFATE
 NADOLOL, NADOLOL
 NAPROXEN SODIUM AND DIPHENHYDRAMINE HYDROCHLORIDE, DIPHENHYDRAMINE HYDROCHLORIDE
 OXAPROZIN, OXAPROZIN
 PARICALCITOL, PARICALCITOL
 PHYTONADIONE, PHYTONADIONE
 PREGABALIN, PREGABALIN
 SEVELAMER CARBONATE, SEVELAMER CARBONATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

* AMNEAL PHARMACEUTICALS CO GMBH

SILODOSIN, SILODOSIN
 SOLIFENACIN SUCCINATE, SOLIFENACIN SUCCINATE
 TADALAFIL, TADALAFIL
 TIAGABINE HYDROCHLORIDE, TIAGABINE HYDROCHLORIDE
 TRANEXAMIC ACID, TRANEXAMIC ACID
 URSODIOL, URSODIOL

AMNEAL PHARMS LLC

* AMNEAL PHARMACEUTICALS LLC

ACTIVELLA, ESTRADIOL
 AMINOCAPROIC ACID, AMINOCAPROIC ACID
 AZATHIOPRINE, AZATHIOPRINE
 AZELASTINE HYDROCHLORIDE, AZELASTINE HYDROCHLORIDE
 AZITHROMYCIN, AZITHROMYCIN
 CLOBAZAM, CLOBAZAM
 CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
 DEXTROMETHORPHAN POLISTIREX, DEXTROMETHORPHAN POLISTIREX (OTC)
 ELURYNG, ETHINYL ESTRADIOL
 ESTRADIOL, ESTRADIOL
 FENOFIBRATE, FENOFIBRATE
 FLUOCINONIDE, FLUOCINONIDE
 MESALAMINE, MESALAMINE
 NYSTATIN AND TRIAMCINOLONE ACETONIDE, NYSTATIN
 OLOPATADINE HYDROCHLORIDE, OLOPATADINE HYDROCHLORIDE
 POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
 PYRIDOSTIGMINE BROMIDE, PYRIDOSTIGMINE BROMIDE
 RITONAVIR, RITONAVIR
 SIROLIMUS, SIROLIMUS
 SUCRALFATE, SUCRALFATE
 TESTOSTERONE, TESTOSTERONE
 TIGECYCLINE, TIGECYCLINE
 TRIENTINE HYDROCHLORIDE, TRIENTINE HYDROCHLORIDE

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
 NAPROXEN SODIUM, NAPROXEN SODIUM (OTC)
 NAPROXEN, NAPROXEN
 OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN
 PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
 REPRESXAIN, HYDROCODONE BITARTRATE
 SULFAMETHOXAZOLE AND TRIMETHOPRIM, SULFAMETHOXAZOLE

* AMNEAL PHARMACEUTICALS OF NY LLC

BEXAROTENE, BEXAROTENE
 CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
 ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM (OTC)
 HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
 ISOTRETINOIN, ISOTRETINOIN
 OSELTAMIVIR PHOSPHATE, OSELTAMIVIR PHOSPHATE
 PROGESTERONE, PROGESTERONE
 SILDENAFIL CITRATE, SILDENAFIL CITRATE
 TETRACYCLINE HYDROCHLORIDE, TETRACYCLINE HYDROCHLORIDE

AMPHASTAR PHARM

* AMPHASTAR PHARMACEUTICAL INC

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

* AMPHASTAR PHARMACEUTICAL INC
 AMPHADASE, HYALURONIDASE
 ENOXAPARIN SODIUM (PRESERVATIVE FREE), ENOXAPARIN SODIUM
 KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE

AMPHASTAR PHARMS INC

* AMPHASTAR PHARMACEUTICALS INC
 CORTROSYN, COSYNTROPIN
 ENOXAPARIN SODIUM, ENOXAPARIN SODIUM
 ISOPROTERENOL HYDROCHLORIDE, ISOPROTERENOL HYDROCHLORIDE
 MEDROXYPROGESTERONE ACETATE, MEDROXYPROGESTERONE ACETATE
 NEOSTIGMINE METHYLSULFATE, NEOSTIGMINE METHYLSULFATE
 SODIUM NITROPRUSSIDE, SODIUM NITROPRUSSIDE

AMRING PHARMS

* AMRING PHARMACEUTICALS INC
 ARSENIC TRIOXIDE, ARSENIC TRIOXIDE
 LATANOPROST, LATANOPROST
 MESALAMINE, MESALAMINE
 NEOMYCIN AND POLYMYXIN B SULFATES AND GRAMICIDIN, GRAMICIDIN
 NEOMYCIN AND POLYMYXIN B SULFATES AND HYDROCORTISONE, HYDROCORTISONE
 NEOSTIGMINE METHYLSULFATE, NEOSTIGMINE METHYLSULFATE
 SUCCINYLCHOLINE CHLORIDE, SUCCINYLCHOLINE CHLORIDE

AMTA

* AMTA LABS LTD
 PROPRANOLOL HYDROCHLORIDE, PROPRANOLOL HYDROCHLORIDE

ANACOR PHARMS INC

* ANACOR PHARMACEUTICALS INC
 EUCRISA, CRISABOROLE
 KERYDIN, TAVABOROLE

ANBEX

* ANBEX INC
 IOSAT, POTASSIUM IODIDE (OTC)

ANBISON LAB

* ANBISON LABORATORY CO LTD
 BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
 MONTELUKAST SODIUM, MONTELUKAST SODIUM
 TAMSULOSIN HYDROCHLORIDE, TAMSULOSIN HYDROCHLORIDE

ANCHEN PHARMS

* ANCHEN PHARMACEUTICALS INC
 ALISKIREN HEMIFUMARATE, ALISKIREN HEMIFUMARATE
 BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
 CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
 DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE
 DUTASTERIDE AND TAMSULOSIN HYDROCHLORIDE, DUTASTERIDE
 FENOFIBRIC ACID, CHOLINE FENOFIBRATE
 FLUVOXAMINE MALEATE, FLUVOXAMINE MALEATE
 LAMOTRIGINE, LAMOTRIGINE
 LEVETIRACETAM, LEVETIRACETAM
 MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
 POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
 PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE
 QUETIAPINE FUMARATE, QUETIAPINE FUMARATE
 TAMSULOSIN HYDROCHLORIDE, TAMSULOSIN HYDROCHLORIDE
 TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE
 TRETINOIN, TRETINOIN
 ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE

* ANCHEN PHARMACEUTICALS TAIWAN INC
 DIVALPROEX SODIUM, DIVALPROEX SODIUM

* ANCHEN PHARMACEUTICALS, INC
 ALPRAZOLAM, ALPRAZOLAM
 CIPROFLOXACIN EXTENDED RELEASE, CIPROFLOXACIN

ANDA REPOSITORY

* ANDA REPOSITORY LLC
 ACETAMINOPHEN AND CODEINE PHOSPHATE, ACETAMINOPHEN
 AMANTADINE HYDROCHLORIDE, AMANTADINE HYDROCHLORIDE
 BENZPHETAMINE HYDROCHLORIDE, BENZPHETAMINE HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ******* ANDA REPOSITORY LLC**

BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
 CICLOPIROX, CICLOPIROX
 CYCLOBENZAPRINE HYDROCHLORIDE, CYCLOBENZAPRINE HYDROCHLORIDE
 CYPROHEPTADINE HYDROCHLORIDE, CYPROHEPTADINE HYDROCHLORIDE
 FLAC, FLUOCINOLONE ACETONIDE
 FLUTICASONE PROPIONATE, FLUTICASONE PROPIONATE
 HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
 IMIQUIMOD, IMIQUIMOD
 ISONIAZID, ISONIAZID
 LAMIVUDINE AND ZIDOVUDINE, LAMIVUDINE
 LEVETIRACETAM, LEVETIRACETAM
 MOMETASONE FUROATE, MOMETASONE FUROATE
 ORPHENADRINE CITRATE, ORPHENADRINE CITRATE
 OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN
 PRIMIDONE, PRIMIDONE
 RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE

ANDOR PHARMS*** ANDOR PHARMACEUTICALS LLC**

METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE

ANI PHARMS*** ANI PHARMACEUTICALS INC**

CORTENEMA, HYDROCORTISONE
 LUVOX, FLUVOXAMINE MALEATE
 METOCLOPRAMIDE HYDROCHLORIDE, METOCLOPRAMIDE HYDROCHLORIDE
 REGLAN, METOCLOPRAMIDE HYDROCHLORIDE

ANI PHARMS INC*** ANI PHARMACEUTICALS INC**

ARIMIDEX, ANASTROZOLE
 ATACAND HCT, CANDESARTAN CILEXETIL
 ATACAND, CANDESARTAN CILEXETIL
 BEXAROTENE, BEXAROTENE
 BRETHINE, TERBUTALINE SULFATE
 CARBIDOPA, CARBIDOPA
 CASODEX, BICALUTAMIDE
 CHOLESTYRAMINE, CHOLESTYRAMINE
 CYCLOPHOSPHAMIDE, CYCLOPHOSPHAMIDE
 DESIPRAMINE HYDROCHLORIDE, DESIPRAMINE HYDROCHLORIDE
 DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE
 DIPHENOXYLATE HYDROCHLORIDE AND ATROPINE SULFATE, ATROPINE SULFATE
 ERYTHROMYCIN ETHYLSUCCINATE, ERYTHROMYCIN ETHYLSUCCINATE
 ETODOLAC, ETODOLAC
 EZETIMIBE AND SIMVASTATIN, EZETIMIBE
 FELBAMATE, FELBAMATE
 FENOFIBRATE (MICRONIZED), FENOFIBRATE
 FLECAINIDE ACETATE, FLECAINIDE ACETATE
 INDAPAMIDE, INDAPAMIDE
 INDERAL LA, PROPRANOLOL HYDROCHLORIDE
 INNOPRAN XL, PROPRANOLOL HYDROCHLORIDE
 LITHOBID, LITHIUM CARBONATE
 MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
 METHAZOLAMIDE, METHAZOLAMIDE
 METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
 MORPHINE SULFATE, MORPHINE SULFATE
 NILUTAMIDE, NILUTAMIDE
 OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
 PENICILLAMINE, PENICILLAMINE
 PINDOLOL, PINDOLOL
 POLYETHYLENE GLYCOL 3350, POLYETHYLENE GLYCOL 3350 (OTC)
 POTASSIUM CITRATE, POTASSIUM CITRATE
 PROPAFENONE HYDROCHLORIDE, PROPAFENONE HYDROCHLORIDE
 RANOLAZINE, RANOLAZINE
 SULFAMETHOXAZOLE AND TRIMETHOPRIM, SULFAMETHOXAZOLE
 VANCOCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE

ANIMA

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

* ANIMA PHARMACEUTICALS PVT LTD
 BETAMETHASONE VALERATE, BETAMETHASONE VALERATE
 DEXAMETHASONE, DEXAMETHASONE

ANNORA

* ANNORA PHARMA PRIVATE LTD
 LAMIVUDINE, LAMIVUDINE

ANNORA PHARMA

* ANNORA PHARMA PRIVATE LTD
 VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE

ANTARES PHARMA INC

* ANTARES PHARMA INC
 OTREXUP, METHOTREXATE
 SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
 XYOSTED (AUTOINJECTOR), TESTOSTERONE ENANTHATE

ANTIBIOTICE

* ANTIBIOTICE SA
 AMPICILLIN AND SULBACTAM, AMPICILLIN SODIUM
 AMPICILLIN SODIUM, AMPICILLIN SODIUM
 NAFCILLIN SODIUM, NAFCILLIN SODIUM

ANTRIM PHARMS LLC

* ANTRIM PHARMACEUTICALS LLC
 ESCITALOPRAM OXALATE, ESCITALOPRAM OXALATE

APEX PHARMS INC

* APEX PHARMACEUTICALS INC
 CYPROHEPTADINE HYDROCHLORIDE, CYPROHEPTADINE HYDROCHLORIDE

APGDI

* ASTELLAS PHARMA GLOBAL DEVELOPMENT INC
 MYRBETRIQ, MIRABEGRON

APIL

* ALLERGAN PHARMACEUTICALS INTERNATIONAL LTD
 ACTONEL, RISEDRONATE SODIUM
 ASACOL HD, MESALAMINE
 ATELVIA, RISEDRONATE SODIUM
 DELZICOL, MESALAMINE
 ENABLEX, DARIFENACIN HYDROBROMIDE
 ESTROSTEP FE, ETHINYL ESTRADIOL
 FEMHRT, ETHINYL ESTRADIOL
 LO LOESTRIN FE, ETHINYL ESTRADIOL
 MINASTRIN 24 FE, ETHINYL ESTRADIOL
 NORCO, ACETAMINOPHEN
 NORETHINDRONE AND ETHINYL ESTRADIOL AND FERROUS FUMARATE, ETHINYL ESTRADIOL
 SARAFEM, FLUOXETINE HYDROCHLORIDE
 TAYTULLA, ETHINYL ESTRADIOL

APNAR PHARMA LP

* APNAR PHARMA LP
 BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
 FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 PRAVASTATIN SODIUM, PRAVASTATIN SODIUM
 TERAZOSIN HYDROCHLORIDE, TERAZOSIN HYDROCHLORIDE

APOLLO

* APOLLO PHARMACEUTICALS INC
 DIHYDROERGOTAMINE MESYLATE, DIHYDROERGOTAMINE MESYLATE
 PIPERACILLIN AND TAZOBACTAM, PIPERACILLIN SODIUM
 TRANEXAMIC ACID, TRANEXAMIC ACID

APOPHARMA INC

* APOPHARMA INC
 FERRIPROX, DEFERIPRONE

APOTEX

* APOTEX INC
 ABIRATERONE ACETATE, ABIRATERONE ACETATE
 ACYCLOVIR, ACYCLOVIR
 ADEFOVIR DIPIVOXIL, ADEFOVIR DIPIVOXIL
 ALENDRONATE SODIUM, ALENDRONATE SODIUM
 ALKERAN, MELPHALAN

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

**** A ****

* APOTEX INC
 ALKERAN, MELPHALAN HYDROCHLORIDE
 AMLODIPINE BESYLATE AND ATORVASTATIN CALCIUM, AMLODIPINE BESYLATE
 AMLODIPINE BESYLATE AND BENAZEPRIL HYDROCHLORIDE, AMLODIPINE BESYLATE
 AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
 ATOMOXETINE HYDROCHLORIDE, ATOMOXETINE HYDROCHLORIDE
 ATOVAQUONE, ATOVAQUONE
 AZELASTINE HYDROCHLORIDE AND FLUTICASONE PROPIONATE, AZELASTINE HYDROCHLORIDE
 BENAZEPRIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, BENAZEPRIL HYDROCHLORIDE
 BUSULFAN, BUSULFAN
 CARBIDOPA AND LEVODOPA, CARBIDOPA
 CARFILZOMIB, CARFILZOMIB
 CEFUROXIME AXETIL, CEFUROXIME AXETIL
 CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 CIMETIDINE, CIMETIDINE (OTC)
 CYCLOBENZAPRINE HYDROCHLORIDE, CYCLOBENZAPRINE HYDROCHLORIDE
 CYCLOSPORINE, CYCLOSPORINE
 DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
 DIVALPROEX SODIUM, DIVALPROEX SODIUM
 DOXAZOSIN MESYLATE, DOXAZOSIN MESYLATE
 DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
 DROSPIRENONE AND ETHINYL ESTRADIOL, DROSPIRENONE
 DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE
 EPINASTINE HYDROCHLORIDE, EPINASTINE HYDROCHLORIDE
 ERLOTINIB HYDROCHLORIDE, ERLOTINIB HYDROCHLORIDE
 ETODOLAC, ETODOLAC
 EZETIMIBE, EZETIMIBE
 FAMCICLOVIR, FAMCICLOVIR
 FAMOTIDINE, FAMOTIDINE
 FENOFIBRATE (MICRONIZED), FENOFIBRATE
 FLUTICASONE PROPIONATE, FLUTICASONE PROPIONATE (OTC)
 FLUVOXAMINE MALEATE, FLUVOXAMINE MALEATE
 FOSAPREPITANT DIMEGLUMINE, FOSAPREPITANT DIMEGLUMINE
 GEMFIBROZIL, GEMFIBROZIL
 GLIPIZIDE, GLIPIZIDE
 GLYCOPYRROLATE, GLYCOPYRROLATE
 GUANFACINE HYDROCHLORIDE, GUANFACINE HYDROCHLORIDE
 IBANDRONATE SODIUM, IBANDRONATE SODIUM
 IMATINIB MESYLATE, IMATINIB MESYLATE
 LAMIVUDINE, LAMIVUDINE
 LEVETIRACETAM, LEVETIRACETAM
 MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 MIDODRINE HYDROCHLORIDE, MIDODRINE HYDROCHLORIDE
 MOXIFLOXACIN HYDROCHLORIDE, MOXIFLOXACIN HYDROCHLORIDE
 NALTREXONE HYDROCHLORIDE, NALTREXONE HYDROCHLORIDE
 NAPROXEN SODIUM AND DIPHENHYDRAMINE HYDROCHLORIDE, DIPHENHYDRAMINE HYDROCHLORIDE
 OLOPATADINE HYDROCHLORIDE, OLOPATADINE HYDROCHLORIDE
 OMEGA-3-ACID ETHYL ESTERS, OMEGA-3-ACID ETHYL ESTERS
 OMEPRAZOLE, OMEPRAZOLE
 ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
 PAROXETINE HYDROCHLORIDE, PAROXETINE HYDROCHLORIDE
 PENTOXIFYLLINE, PENTOXIFYLLINE
 POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
 PRAVASTATIN SODIUM, PRAVASTATIN SODIUM
 RAMIPRIL, RAMIPRIL
 RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
 SELEGILINE HYDROCHLORIDE, SELEGILINE HYDROCHLORIDE
 SIROLIMUS, SIROLIMUS
 SOTALOL HYDROCHLORIDE, SOTALOL HYDROCHLORIDE
 TAMOXIFEN CITRATE, TAMOXIFEN CITRATE
 TETRABENZAZINE, TETRABENZAZINE
 TICLOPIDINE HYDROCHLORIDE, TICLOPIDINE HYDROCHLORIDE
 TIGECYCLINE, TIGECYCLINE
 TIMOLOL MALEATE, TIMOLOL MALEATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ******* APOTEX INC**

TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE
 TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE
 TRANEXAMIC ACID, TRANEXAMIC ACID
 TRAVOPROST, TRAVOPROST
 TRAZODONE HYDROCHLORIDE, TRAZODONE HYDROCHLORIDE
 TROSPIUM CHLORIDE, TROSPIUM CHLORIDE
 VALACYCLOVIR HYDROCHLORIDE, VALACYCLOVIR HYDROCHLORIDE
 ZIPRASIDONE HYDROCHLORIDE, ZIPRASIDONE HYDROCHLORIDE
 ZOLEDRONIC ACID, ZOLEDRONIC ACID

APOTEX CORP*** APOTEX CORP**

QUINAPRIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 RILUZOLE, RILUZOLE

APOTEX INC*** APOTEX INC**

ABACAVIR SULFATE, ABACAVIR SULFATE
 ALFUZOSIN HYDROCHLORIDE, ALFUZOSIN HYDROCHLORIDE
 ALPRAZOLAM, ALPRAZOLAM
 ARIPIPIRAZOLE, ARIPIPIRAZOLE
 ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM
 AZELASTINE HYDROCHLORIDE, AZELASTINE HYDROCHLORIDE
 BENZONATATE, BENZONATATE
 BICALUTAMIDE, BICALUTAMIDE
 BIMATOPROST, BIMATOPROST
 BUTORPHANOL TARTRATE, BUTORPHANOL TARTRATE
 CALCITONIN-SALMON, CALCITONIN SALMON
 CARBAMAZEPINE, CARBAMAZEPINE
 CARBIDOPA AND LEVODOPA, CARBIDOPA
 CEFPROZIL, CEFPROZIL
 CELECOXIB, CELECOXIB
 CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 CLOPIDOGREL BISULFATE, CLOPIDOGREL BISULFATE
 DICLOFENAC SODIUM, DICLOFENAC SODIUM
 ENOXAPARIN SODIUM (PRESERVATIVE FREE), ENOXAPARIN SODIUM
 GRANISETRON HYDROCHLORIDE, GRANISETRON HYDROCHLORIDE
 IBANDRONATE SODIUM, IBANDRONATE SODIUM
 IMIQUIMOD, IMIQUIMOD
 IPRATROPIUM BROMIDE, IPRATROPIUM BROMIDE
 KETOTIFEN FUMARATE, KETOTIFEN FUMARATE (OTC)
 LETROZOLE, LETROZOLE
 LEVETIRACETAM, LEVETIRACETAM
 LEVOFLOXACIN, LEVOFLOXACIN
 MODAFINIL, MODAFINIL
 MOEXIPRIL HYDROCHLORIDE, MOEXIPRIL HYDROCHLORIDE
 MOMETASONE FUROATE, MOMETASONE FUROATE
 MYCOPHENOLIC ACID, MYCOPHENOLIC ACID
 OFLOXACIN, OFLOXACIN
 OLANZAPINE, OLANZAPINE
 OLOPATADINE HYDROCHLORIDE, OLOPATADINE HYDROCHLORIDE
 RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE (OTC)
 RISEDRONATE SODIUM, RISEDRONATE SODIUM
 RISPERIDONE, RISPERIDONE
 RIVASTIGMINE TARTRATE, RIVASTIGMINE TARTRATE
 ROSUVASTATIN CALCIUM, ROSUVASTATIN CALCIUM
 SERTRALINE HYDROCHLORIDE, SERTRALINE HYDROCHLORIDE
 TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE
 TRANEXAMIC ACID, TRANEXAMIC ACID
 TRAZODONE HYDROCHLORIDE, TRAZODONE HYDROCHLORIDE
 TRIAMTERENE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE

*** APOTEX INC ETOBICOKE SITE**

ACYCLOVIR, ACYCLOVIR
 ALLOPURINOL, ALLOPURINOL
 BALSALAZIDE DISODIUM, BALSALAZIDE DISODIUM

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

**** A ****

* APOTEX INC ETOBICOKE SITE
 BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
 CARBAMAZEPINE, CARBAMAZEPINE
 CILOSTAZOL, CILOSTAZOL
 CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE
 DESMOPRESSIN ACETATE, DESMOPRESSIN ACETATE
 DILTZAC, DILTIAZEM HYDROCHLORIDE
 ETODOLAC, ETODOLAC
 FOSINOPRIL SODIUM, FOSINOPRIL SODIUM
 GABAPENTIN, GABAPENTIN
 LEFLUNOMIDE, LEFLUNOMIDE
 LORATADINE, LORATADINE (OTC)
 MELOXICAM, MELOXICAM
 MIRTAZAPINE, MIRTAZAPINE
 OXAPROZIN, OXAPROZIN
 SELEGILINE HYDROCHLORIDE, SELEGILINE HYDROCHLORIDE
 TOPIRAMATE, TOPIRAMATE
 ZONISAMIDE, ZONISAMIDE

* APOTEX INC RICHMOND HILL
 AZELASTINE HYDROCHLORIDE, AZELASTINE HYDROCHLORIDE
 BUDESONIDE, BUDESONIDE (OTC)
 DESMOPRESSIN ACETATE (NEEDS NO REFRIGERATION), DESMOPRESSIN ACETATE
 FLUTICASONE PROPIONATE, FLUTICASONE PROPIONATE
 KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE

* APOTEX INC.
 DILTZAC, DILTIAZEM HYDROCHLORIDE
 GALANTAMINE HYDROBROMIDE, GALANTAMINE HYDROBROMIDE

APOTEX TECHNOLOGIES

* APOTEX TECHNOLOGIES INC
 PAXIL CR, PAROXETINE HYDROCHLORIDE
 PAXIL, PAROXETINE HYDROCHLORIDE

APOTHECON

* APOTHECON INC DIV BRISTOL MYERS SQUIBB
 KENALOG-10, TRIAMCINOLONE ACETONIDE
 KENALOG-40, TRIAMCINOLONE ACETONIDE
 KENALOG-80, TRIAMCINOLONE ACETONIDE

APP PHARMS

* APP PHARMACEUTICALS LLC
 DIPHENHYDRAMINE HYDROCHLORIDE, DIPHENHYDRAMINE HYDROCHLORIDE

APPCO

* APPCO PHARMA LLC
 ALBUTEROL SULFATE, ALBUTEROL SULFATE
 BUDESONIDE, BUDESONIDE
 FENOFIBRATE, FENOFIBRATE
 HALOPERIDOL, HALOPERIDOL
 METAXALONE, METAXALONE
 MODAFINIL, MODAFINIL
 RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
 SILDENAFIL CITRATE, SILDENAFIL CITRATE
 VORICONAZOLE, VORICONAZOLE

APRECIA PHARMS

* APRECIA PHARMACEUTICALS LLC
 SPRITAM, LEVETIRACETAM

APTAPHARMA INC

* APTAPHARMA INC
 IBUPROFEN, IBUPROFEN (OTC)

AQUESTIVE THERAP

* AQUESTIVE THERAPEUTICS
 EXSERVAN, RILUZOLE
 SYMPAZAN, CLOBAZAM

ARBOR PHARMS LLC

* ARBOR PHARMACEUTICALS LLC
 BIDIL, HYDRALAZINE HYDROCHLORIDE
 DEXTROAMPHETAMINE SULFATE, DEXTROAMPHETAMINE SULFATE
 E.E.S. 400, ERYTHROMYCIN ETHYLSUCCINATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

**** A ****

* ARBOR PHARMACEUTICALS LLC
 E.E.S., ERYTHROMYCIN ETHYLSUCCINATE
 EDARBI, AZILSARTAN KAMEDOXOMIL
 EDARBYCLOR, AZILSARTAN KAMEDOXOMIL
 ERY-TAB, ERYTHROMYCIN
 ERYPED, ERYTHROMYCIN ETHYLSUCCINATE
 ERYTHROCIN STEARATE, ERYTHROMYCIN STEARATE
 ERYTHROMYCIN ETHYLSUCCINATE, ERYTHROMYCIN ETHYLSUCCINATE
 ERYTHROMYCIN, ERYTHROMYCIN
 EVEKEO ODT, AMPHETAMINE SULFATE
 EVEKEO, AMPHETAMINE SULFATE
 GLIADEL, CARMUSTINE
 HORIZANT, GABAPENTIN ENACARBIL
 NYMALIZE, NIMODIPINE
 SKLICE, IVERMECTIN
 SOTYLIZE, SOTALOL HYDROCHLORIDE
 TRIPTODUR KIT, TRIPTORELIN PAMOATE

ARCO PHARMS LLC

* ARCO PHARMACEUTICALS LLC
 THYROSHIELD, POTASSIUM IODIDE (OTC)

ARDELYX INC

* ARDELYX INC
 IBSRELA, TENAPANOR HYDROCHLORIDE

AREVA PHARMS

* AREVA PHARMACEUTICALS INC
 FLUDARABINE PHOSPHATE, FLUDARABINE PHOSPHATE
 PAMIDRONATE DISODIUM, PAMIDRONATE DISODIUM
 VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE

ARIAD

* ARIAD PHARMACEUTICALS INC
 ALUNBRIG, BRIGATINIB
 ICLUSIG, PONATINIB HYDROCHLORIDE

ARISE

* ARISE PHARMACEUTICALS LLC
 ALBUTEROL SULFATE, ALBUTEROL SULFATE
 CHLORTHALIDONE, CHLORTHALIDONE
 IBUPROFEN, IBUPROFEN (OTC)
 LAMIVUDINE, LAMIVUDINE
 OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE

ARMSTRONG PHARMS

* ARMSTRONG PHARMACEUTICALS INC
 PRIMATENE MIST, EPINEPHRINE (OTC)

ARRAY BIOPHARMA INC

* ARRAY BIOPHARMA INC
 BRAFTOVI, ENCORAFENIB
 MEKTOVI, BINIMETINIB

ASCEND THERAPS US

* ASCEND THERAPEUTICS US LLC
 BINOSTO, ALENDRONATE SODIUM
 ESTROGEL, ESTRADIOL

ASCENT PHARMS INC

* ASCENT PHARMACEUTICALS INC
 BENZONATATE, BENZONATATE
 DRONABINOL, DRONABINOL
 DUTASTERIDE, DUTASTERIDE
 HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
 HYDROMORPHONE HYDROCHLORIDE, HYDROMORPHONE HYDROCHLORIDE
 IBUPROFEN, IBUPROFEN (OTC)
 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

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ******ASPEN**

* ASPEN PHARMA USA INC
HYDROXYPROGESTERONE CAPROATE, HYDROXYPROGESTERONE CAPROATE

ASPEN GLOBAL

* ASPEN GLOBAL INC
MYLERAN, BUSULFAN

ASPEN GLOBAL INC

* ASPEN GLOBAL INC
BENZTROPINE MESYLATE, BENZTROPINE MESYLATE
CYCLESSA, DESOGESTREL
HYDROXYPROGESTERONE CAPROATE, HYDROXYPROGESTERONE CAPROATE
LEUKERAN, CHLORAMBUCIL
THIOGUANINE, THIOGUANINE

ASSERTIO

* ASSERTIO THERAPEUTICS INC
CAMBIA, DICLOFENAC POTASSIUM
ZIPSOR, DICLOFENAC POTASSIUM

ASTELLAS

* ASTELLAS PHARMA US INC
AMBISOME, AMPHOTERICIN B
ASTAGRAF XL, TACROLIMUS
CRESEMBA, ISAVUCONAZONIUM SULFATE
LEXISCAN, REGADENOSON
MYCAMINE, MICAFUNGIN SODIUM
PROGRAF, TACROLIMUS
VESICARE, SOLIFENACIN SUCCINATE
XOSPATA, GILTERITINIB FUMARATE
XTANDI, ENZALUTAMIDE

ASTRAL

* ASTRAL STERITECH PVT LTD
AMPICILLIN AND SULBACTAM, AMPICILLIN SODIUM
CEFTRIAZONE SODIUM, CEFTRIAZONE SODIUM
CEFTRIAZONE, CEFTRIAZONE SODIUM

ASTRAZENECA

* ASTRAZENECA LP
PULMICORT FLEXHALER, BUDESONIDE
SYMBICORT, BUDESONIDE
* ASTRAZENECA PHARMACEUTICALS LP
FASLODEX, FULVESTRANT
ZOMIG, ZOLMITRIPTAN
ZOMIG-ZMT, ZOLMITRIPTAN
* ASTRAZENECA UK LTD
CALQUENCE, ACALABRUTINIB
SEROQUEL XR, QUETIAPINE FUMARATE

ASTRAZENECA AB

* ASTRAZENECA AB
BYDUREON BCISE, EXENATIDE
BYDUREON PEN, EXENATIDE SYNTHETIC
BYDUREON, EXENATIDE SYNTHETIC
BYETTA, EXENATIDE SYNTHETIC
FARXIGA, DAPAGLIFLOZIN
KOMBIGLYZE XR, METFORMIN HYDROCHLORIDE
ONGLYZA, SAXAGLIPTIN HYDROCHLORIDE
QTERN, DAPAGLIFLOZIN
QTERNMET XR, DAPAGLIFLOZIN
SYMLIN, PRAMLINTIDE ACETATE
XIGDUO XR, DAPAGLIFLOZIN

ASTRAZENECA LP

* ASTRAZENECA LP
NEXIUM 24HR, ESOMEPRAZOLE MAGNESIUM (OTC)

ASTRAZENECA PHARMS

* ASTRAZENECA PHARMACEUTICALS LP
BEVESPI AEROSPHERE, FORMOTEROL FUMARATE
BRILINTA, TICAGRELOR

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ******* ASTRAZENECA PHARMACEUTICALS LP**

DALIRESP, ROFLUMILAST
 IRESSA, GEFITINIB
 LOKELMA, SODIUM ZIRCONIUM CYCLOSILICATE
 LYNPARZA, OLAPARIB
 MOVANTIK, NALOXEGOL OXALATE
 NEXIUM IV, ESOMEPRAZOLE SODIUM
 NEXIUM, ESOMEPRAZOLE MAGNESIUM
 PRILOSEC OTC, OMEPRAZOLE MAGNESIUM (OTC)
 PULMICORT RESPULES, BUDESONIDE
 RHINOCORT ALLERGY, BUDESONIDE (OTC)
 SEROQUEL, QUETIAPINE FUMARATE
 TAGRISSO, OSIMERTINIB MESYLATE

ATHEM*** ATHEM HOLDINGS LLC**

CHILDREN'S CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 CHILDREN'S CETIRIZINE HYDROCHLORIDE HIVES RELIEF, CETIRIZINE HYDROCHLORIDE (OTC)
 FOLIC ACID, FOLIC ACID
 LABETALOL HYDROCHLORIDE, LABETALOL HYDROCHLORIDE
 OXYBUTYNYN CHLORIDE, OXYBUTYNYN CHLORIDE

ATHENA*** ATHENA BIOSCIENCES LLC**

FIBRICOR, FENOFIBRIC ACID

ATHENEX*** ATHENEX PHARMACEUTICAL DIV**

PENTOBARBITAL SODIUM, PENTOBARBITAL SODIUM

ATHENEX INC*** ATHENEX INC**

BUSULFAN, BUSULFAN
 DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
 DIPYRIDAMOLE, DIPYRIDAMOLE
 DOXAPRAM HYDROCHLORIDE, DOXAPRAM HYDROCHLORIDE
 ENALAPRILAT, ENALAPRILAT
 FAMOTIDINE PRESERVATIVE FREE, FAMOTIDINE
 FAMOTIDINE, FAMOTIDINE
 PROCHLORPERAZINE EDISYLATE, PROCHLORPERAZINE EDISYLATE
 PROPRANOLOL HYDROCHLORIDE, PROPRANOLOL HYDROCHLORIDE
 TERBUTALINE SULFATE, TERBUTALINE SULFATE
 VALPROATE SODIUM, VALPROATE SODIUM

ATLAS PHARMS LLC*** ATLAS PHARMACEUTICALS LLC**

METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE

ATNAHS PHARMA US*** ATNAHS PHARMA US LTD**

ANAPROX DS, NAPROXEN SODIUM
 EC-NAPROSYN, NAPROXEN
 NAPROSYN, NAPROXEN

ATON*** ATON PHARMA INC**

LODOSYN, CARBIDOPA

AUCTA*** AUCTA PHARMACEUTICALS INC**

VIGADRONE, VIGABATRIN

AUREX*** AUREX LABORATORIES LTD LIABILITY CO**

MECLIZINE HYDROCHLORIDE, MECLIZINE HYDROCHLORIDE

AUROBINDO*** AUROBINDO PHARMA LTD**

AMOXICILLIN, AMOXICILLIN
 CEFADROXIL, CEFADROXIL/CEFADROXIL HEMIHYDRATE
 CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE
 CLARITHROMYCIN, CLARITHROMYCIN
 DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
 LISINAPRIL AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ******* AUROBINDO PHARMA LTD**

LISINOPRIL, LISINOPRIL
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 MIRTAZAPINE, MIRTAZAPINE
 NEVIRAPINE, NEVIRAPINE
 ZIDOVUDINE, ZIDOVUDINE

AUROBINDO PHARMA*** AUROBINDO PHARMA**

AMPICILLIN AND SULBACTAM, AMPICILLIN SODIUM

*** AUROBINDO PHARMA LTD**

ALENDRONATE SODIUM, ALENDRONATE SODIUM
 AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
 AMPICILLIN SODIUM, AMPICILLIN SODIUM
 ATENOLOL, ATENOLOL
 BENAZEPRIL HYDROCHLORIDE, BENAZEPRIL HYDROCHLORIDE
 BISOPROLOL FUMARATE, BISOPROLOL FUMARATE
 CARISOPRODOL, CARISOPRODOL
 CARVEDILOL, CARVEDILOL
 CEFADROXIL, CEFADROXIL/CEFADROXIL HEMIHYDRATE
 CEFDINIR, CEFDINIR
 CEFPODOXIME PROXETIL, CEFPODOXIME PROXETIL
 CEFPROZIL, CEFPROZIL
 CHILDREN'S CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 CHILDREN'S CETIRIZINE HYDROCHLORIDE HIVES RELIEF, CETIRIZINE HYDROCHLORIDE (OTC)
 CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE
 CLINDAMYCIN HYDROCHLORIDE, CLINDAMYCIN HYDROCHLORIDE
 CYCLOBENZAPRINE HYDROCHLORIDE, CYCLOBENZAPRINE HYDROCHLORIDE
 DIDANOSINE, DIDANOSINE
 FINASTERIDE, FINASTERIDE
 FLUCONAZOLE, FLUCONAZOLE
 FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 FOSINOPRIL SODIUM AND HYDROCHLOROTHIAZIDE, FOSINOPRIL SODIUM
 GLYBURIDE AND METFORMIN HYDROCHLORIDE, GLYBURIDE
 GLYBURIDE, GLYBURIDE
 HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 LAMOTRIGINE, LAMOTRIGINE
 LEVETIRACETAM, LEVETIRACETAM
 LOSARTAN POTASSIUM AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 LOSARTAN POTASSIUM, LOSARTAN POTASSIUM
 MELOXICAM, MELOXICAM
 METOPROLOL TARTRATE, METOPROLOL TARTRATE
 MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE
 ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
 ONDANSETRON, ONDANSETRON
 PAROXETINE HYDROCHLORIDE, PAROXETINE HYDROCHLORIDE
 PENICILLIN V POTASSIUM, PENICILLIN V POTASSIUM
 PERINDOPRIL ERBUMINE, PERINDOPRIL ERBUMINE
 QUINAPRIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 RIBAVIRIN, RIBAVIRIN
 SERTRALINE HYDROCHLORIDE, SERTRALINE HYDROCHLORIDE
 SIMVASTATIN, SIMVASTATIN
 STAVUDINE, STAVUDINE
 SULFAMETHOXAZOLE AND TRIMETHOPRIM, SULFAMETHOXAZOLE
 SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
 TERBINAFINE HYDROCHLORIDE, TERBINAFINE HYDROCHLORIDE
 TOPIRAMATE, TOPIRAMATE
 TORSEMIDE, TORSEMIDE
 TRANDOLAPRIL, TRANDOLAPRIL
 VALACYCLOVIR HYDROCHLORIDE, VALACYCLOVIR HYDROCHLORIDE
 VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE
 ZALEPLON, ZALEPLON
 ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE

AUROBINDO PHARMA LTD*** AUROBINDO PHARMA LIMITED**

DIVALPROEX SODIUM, DIVALPROEX SODIUM

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

* AUROBINDO PHARMA LIMITED
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, ACETAMINOPHEN (OTC)
ACETYLCYSTEINE, ACETYLCYSTEINE
ACYCLOVIR SODIUM, ACYCLOVIR SODIUM
ADENOSINE, ADENOSINE
AFIRMELLE, ETHINYL ESTRADIOL
ALBUTEROL SULFATE, ALBUTEROL SULFATE
ALFUZOSIN HYDROCHLORIDE, ALFUZOSIN HYDROCHLORIDE
ALPRAZOLAM, ALPRAZOLAM
AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE
AMLODIPINE AND OLMESARTAN MEDOXOMIL, AMLODIPINE BESYLATE
AMLODIPINE BESYLATE AND BENAZEPRIL HYDROCHLORIDE, AMLODIPINE BESYLATE
AMLODIPINE BESYLATE AND VALSARTAN, AMLODIPINE BESYLATE
AMLODIPINE BESYLATE, VALSARTAN AND HYDROCHLOROTHIAZIDE, AMLODIPINE BESYLATE
AMOXICILLIN AND CLAVULANATE POTASSIUM, AMOXICILLIN
AMOXICILLIN, AMOXICILLIN
ARGATROBAN IN SODIUM CHLORIDE, ARGATROBAN
ARIPIPIRAZOLE, ARIPIPIRAZOLE
ARMODAFINIL, ARMODAFINIL
ATAZANAVIR SULFATE, ATAZANAVIR SULFATE
ATHENTIA NEXT, LEVONORGESTREL (OTC)
ATOMOXETINE HYDROCHLORIDE, ATOMOXETINE HYDROCHLORIDE
ATRACURIUM BESYLATE PRESERVATIVE FREE, ATRACURIUM BESYLATE
ATRACURIUM BESYLATE, ATRACURIUM BESYLATE
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
AZITHROMYCIN, AZITHROMYCIN
BIVALIRUDIN, BIVALIRUDIN
BUPIVACAINE HYDROCHLORIDE PRESERVATIVE FREE, BUPIVACAINE HYDROCHLORIDE
BUPIVACAINE HYDROCHLORIDE, BUPIVACAINE HYDROCHLORIDE
BUSPIRONE HYDROCHLORIDE, BUSPIRONE HYDROCHLORIDE
CAFFEINE CITRATE, CAFFEINE CITRATE
CARBIDOPA, CARBIDOPA
CEFIXIME, CEFIXIME
CEFPODOXIME PROXETIL, CEFPODOXIME PROXETIL
CEFPROZIL, CEFPROZIL
CEFUROXIME AXETIL, CEFUROXIME AXETIL
CELECOXIB, CELECOXIB
CEPHALEXIN, CEPHALEXIN
CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
CETIRIZINE HYDROCHLORIDE HIVES RELIEF, CETIRIZINE HYDROCHLORIDE (OTC)
CINACALCET HYDROCHLORIDE, CINACALCET HYDROCHLORIDE
CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE
CLINDAMYCIN PALMITATE HYDROCHLORIDE, CLINDAMYCIN PALMITATE HYDROCHLORIDE
CLOPIDOGREL BISULFATE, CLOPIDOGREL BISULFATE
CLOZAPINE, CLOZAPINE
CYONANZ, ETHINYL ESTRADIOL
DALFAMPRIDINE, DALFAMPRIDINE
DARIFENACIN HYDROBROMIDE, DARIFENACIN HYDROBROMIDE
DESOGESTREL AND ETHINYL ESTRADIOL, DESOGESTREL
DEXAMETHASONE SODIUM PHOSPHATE, DEXAMETHASONE SODIUM PHOSPHATE
DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE
DIVALPROEX SODIUM, DIVALPROEX SODIUM
DOFETILDE, DOFETILDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

* AUROBINDO PHARMA LTD
DORZOLAMIDE HYDROCHLORIDE AND TIMOLOL MALEATE, DORZOLAMIDE HYDROCHLORIDE
DOXEPIN HYDROCHLORIDE, DOXEPIN HYDROCHLORIDE
DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE
EFAVIRENZ, EFAVIRENZ
EFAVIRENZ, EMTRICITABINE, AND TENOFOVIR DISOPROXIL FUMARATE, EFAVIRENZ
ELETRIPTAN HYDROBROMIDE, ELETRIPTAN HYDROBROMIDE
EMTRICITABINE AND TENOFOVIR DISOPROXIL FUMARATE, EMTRICITABINE
ENTACAPONE, ENTACAPONE
ENTECAVIR, ENTECAVIR
EPTIFIBATIDE, EPTIFIBATIDE
ERTAPENEM SODIUM, ERTAPENEM SODIUM
ESCITALOPRAM OXALATE, ESCITALOPRAM OXALATE
ESMOLOL HYDROCHLORIDE, ESMOLOL HYDROCHLORIDE
ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM
ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM (OTC)
ESOMEPRAZOLE SODIUM, ESOMEPRAZOLE SODIUM
ESZOPICLONE, ESZOPICLONE
ETOMIDATE, ETOMIDATE
EZETIMIBE, EZETIMIBE
FAMCICLOVIR, FAMCICLOVIR
FAMOTIDINE, FAMOTIDINE
FAMOTIDINE, FAMOTIDINE (OTC)
FELODIPINE, FELODIPINE
FENOFIBRATE, FENOFIBRATE
FENOFIBRIC ACID, CHOLINE FENOFIBRATE
FESOTERODINE FUMARATE, FESOTERODINE FUMARATE
FEXOFENADINE HYDROCHLORIDE AND PSEUDOEPHEDRINE HYDROCHLORIDE, FEXOFENADINE
FINASTERIDE, FINASTERIDE
FLECAINIDE ACETATE, FLECAINIDE ACETATE
FLUCONAZOLE, FLUCONAZOLE
FLUPHENAZINE DECANOATE, FLUPHENAZINE DECANOATE
FONDAPARINUX SODIUM, FONDAPARINUX SODIUM
FUROSEMIDE, FUROSEMIDE
GABAPENTIN, GABAPENTIN
GALANTAMINE HYDROBROMIDE, GALANTAMINE HYDROBROMIDE
GEMFIBROZIL, GEMFIBROZIL
GLIMEPIRIDE, GLIMEPIRIDE
GLIPIZIDE, GLIPIZIDE
GLYCOPYRROLATE, GLYCOPYRROLATE
GRANISETRON HYDROCHLORIDE, GRANISETRON HYDROCHLORIDE
GUAIFENESIN AND DEXTROMETHORPHAN HYDROBROMIDE, DEXTROMETHORPHAN HYDROBROMIDE (OTC)
GUAIFENESIN, GUAIFENESIN (OTC)
IBANDRONATE SODIUM, IBANDRONATE SODIUM
IBUPROFEN AND DIPHENHYDRAMINE HYDROCHLORIDE, DIPHENHYDRAMINE HYDROCHLORIDE (OTC)
IBUPROFEN AND PSEUDOEPHEDRINE HYDROCHLORIDE, IBUPROFEN (OTC)
IBUPROFEN, IBUPROFEN
IBUPROFEN, IBUPROFEN (OTC)
ICLEVIA, ETHINYL ESTRADIOL
INCASSIA, NORETHINDRONE
INDOMETHACIN, INDOMETHACIN
IPRATROPIUM BROMIDE, IPRATROPIUM BROMIDE
IRBESARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
IRBESARTAN, IRBESARTAN
ISOSULFAN BLUE, ISOSULFAN BLUE
KALLIGA, DESOGESTREL
LAMIVUDINE AND ZIDOVUDINE, LAMIVUDINE
LAMIVUDINE, LAMIVUDINE
LATANOPROST, LATANOPROST
LEVALBUTEROL HYDROCHLORIDE, LEVALBUTEROL HYDROCHLORIDE
LEVETIRACETAM IN SODIUM CHLORIDE, LEVETIRACETAM
LEVETIRACETAM, LEVETIRACETAM
LEVOFLOXACIN IN DEXTROSE 5% IN PLASTIC CONTAINER, LEVOFLOXACIN
LEVOFLOXACIN, LEVOFLOXACIN
LIDOCAINE HYDROCHLORIDE PRESERVATIVE FREE, LIDOCAINE HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

* AUROBINDO PHARMA LTD
LIDOCAINE HYDROCHLORIDE, LIDOCAINE HYDROCHLORIDE
LINEZOLID, LINEZOLID
LO SIMPESSE, ETHINYL ESTRADIOL
LO-ZUMANDIMINE, DROSPIRENONE
LOPERAMIDE HYDROCHLORIDE, LOPERAMIDE HYDROCHLORIDE (OTC)
LORATADINE, LORATADINE (OTC)
MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
MEROPENEM, MEROPENEM
METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
METHENAMINE HIPPURATE, METHENAMINE HIPPURATE
METHOCARBAMOL, METHOCARBAMOL
METHYLPREDNISOLONE SODIUM SUCCINATE, METHYLPREDNISOLONE SODIUM SUCCINATE
METRONIDAZOLE, METRONIDAZOLE
MILI, ETHINYL ESTRADIOL
MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE
MIRTAZAPINE, MIRTAZAPINE
MODAFINIL, MODAFINIL
MONTELUKAST SODIUM, MONTELUKAST SODIUM
MOXIFLOXACIN HYDROCHLORIDE, MOXIFLOXACIN HYDROCHLORIDE
NADOLOL, NADOLOL
NAFCILLIN SODIUM, NAFCILLIN SODIUM
NALOXONE HYDROCHLORIDE, NALOXONE HYDROCHLORIDE
NAPROXEN SODIUM, NAPROXEN SODIUM
NAPROXEN SODIUM, NAPROXEN SODIUM (OTC)
NAPROXEN, NAPROXEN
NEVIRAPINE, NEVIRAPINE
NEXESTA FE, ETHINYL ESTRADIOL
NIACIN, NIACIN
NYLIA 1/35, ETHINYL ESTRADIOL
NYLIA 7/7/7, ETHINYL ESTRADIOL
OLANZAPINE, OLANZAPINE
OLMESARTAN MEDOXOMIL AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
OLMESARTAN MEDOXOMIL, OLMESARTAN MEDOXOMIL
OLOPATADINE HYDROCHLORIDE, OLOPATADINE HYDROCHLORIDE
OMEPRAZOLE MAGNESIUM, OMEPRAZOLE MAGNESIUM (OTC)
OMEPRAZOLE, OMEPRAZOLE
ONDANSETRON HYDROCHLORIDE PRESERVATIVE FREE, ONDANSETRON HYDROCHLORIDE
ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
OXACILLIN SODIUM, OXACILLIN SODIUM
PALONOSETRON HYDROCHLORIDE, PALONOSETRON HYDROCHLORIDE
PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
PARICALCITOL, PARICALCITOL
PHENYTOIN SODIUM, PHENYTOIN SODIUM
PIOGLITAZONE HYDROCHLORIDE AND METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
PIOGLITAZONE HYDROCHLORIDE, PIOGLITAZONE HYDROCHLORIDE
PIPERACILLIN AND TAZOBACTAM, PIPERACILLIN SODIUM
PITAVASTATIN CALCIUM, PITAVASTATIN CALCIUM
POLYETHYLENE GLYCOL 3350, POLYETHYLENE GLYCOL 3350 (OTC)
POLYMYXIN B SULFATE, POLYMYXIN B SULFATE
POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE
PRASUGREL, PRASUGREL HYDROCHLORIDE
PRAVASTATIN SODIUM, PRAVASTATIN SODIUM
PROPAFENONE HYDROCHLORIDE, PROPAFENONE HYDROCHLORIDE
PSEUDOEPHEDRINE HYDROCHLORIDE, PSEUDOEPHEDRINE HYDROCHLORIDE (OTC)
QUETIAPINE FUMARATE, QUETIAPINE FUMARATE
QUINAPRIL HYDROCHLORIDE, QUINAPRIL HYDROCHLORIDE
RABEPRAZOLE SODIUM, RABEPRAZOLE SODIUM
RALOXIFENE HYDROCHLORIDE, RALOXIFENE HYDROCHLORIDE
RAMIPRIL, RAMIPRIL
RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE (OTC)
REPAGLINIDE, REPAGLINIDE
RISEDRONATE SODIUM, RISEDRONATE SODIUM

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

- * AUROBINDO PHARMA LTD
 RISPERIDONE, RISPERIDONE
 RITONAVIR, RITONAVIR
 RIVASTIGMINE TARTRATE, RIVASTIGMINE TARTRATE
 RIZATRIPTAN BENZOATE, RIZATRIPTAN BENZOATE
 ROCURONIUM BROMIDE, ROCURONIUM BROMIDE
 ROPIVACAINE HYDROCHLORIDE, ROPIVACAINE HYDROCHLORIDE
 ROSUVASTATIN CALCIUM, ROSUVASTATIN CALCIUM
 SEVELAMER CARBONATE, SEVELAMER CARBONATE
 SILDENAFIL CITRATE, SILDENAFIL CITRATE
 SILODOSIN, SILODOSIN
 SIMPESE, ETHINYL ESTRADIOL
 SUMATRIPTAN AND NAPROXEN SODIUM, NAPROXEN SODIUM
 SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
 TADALAFIL, TADALAFIL
 TAMSULOSIN HYDROCHLORIDE, TAMSULOSIN HYDROCHLORIDE
 TELMISARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 TELMISARTAN, TELMISARTAN
 TENOFOVIR DISOPROXIL FUMARATE, TENOFOVIR DISOPROXIL FUMARATE
 TIGECYCLINE, TIGECYCLINE
 TRAMADOL HYDROCHLORIDE AND ACETAMINOPHEN, ACETAMINOPHEN
 TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE
 TRANEXAMIC ACID, TRANEXAMIC ACID
 TRI-LO-MILI, ETHINYL ESTRADIOL
 TRI-MILI, ETHINYL ESTRADIOL
 VALGANCICLOVIR HYDROCHLORIDE, VALGANCICLOVIR HYDROCHLORIDE
 VALSARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 VALSARTAN, VALSARTAN
 VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE
 VECURONIUM BROMIDE, VECURONIUM BROMIDE
 VORICONAZOLE, VORICONAZOLE
 ZIPRASIDONE HYDROCHLORIDE, ZIPRASIDONE HYDROCHLORIDE
 ZOLEDRONIC ACID, ZOLEDRONIC ACID
 ZOLMITRIPTAN, ZOLMITRIPTAN
 ZUMANDIMINE, DROSPIRENONE
- * AUROBINDO PHARMA LTD INC
 ZIDOVUDINE, ZIDOVUDINE

AUROLIFE PHARMA LLC

- * AUROLIFE PHARMA LLC
 ACETAMINOPHEN AND CODEINE PHOSPHATE, ACETAMINOPHEN
 AMPHETAMINE SULFATE, AMPHETAMINE SULFATE
 BUTALBITAL, ACETAMINOPHEN AND CAFFEINE, ACETAMINOPHEN
 CHILDREN'S FEXOFENADINE HYDROCHLORIDE ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
 DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE
 DEXTROAMPHETAMINE SULFATE, DEXTROAMPHETAMINE SULFATE
 DUTASTERIDE, DUTASTERIDE
 FEXOFENADINE HYDROCHLORIDE ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
 GLYCOPYRROLATE, GLYCOPYRROLATE
 HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
 HYDROCODONE BITARTRATE AND IBUPROFEN, HYDROCODONE BITARTRATE
 HYDROMORPHONE HYDROCHLORIDE, HYDROMORPHONE HYDROCHLORIDE
 LORAZEPAM, LORAZEPAM
 METHADONE HYDROCHLORIDE, METHADONE HYDROCHLORIDE
 OMEPRAZOLE AND SODIUM BICARBONATE, OMEPRAZOLE
 OMEPRAZOLE AND SODIUM BICARBONATE, OMEPRAZOLE (OTC)
 OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN
 OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
 OXYMORPHONE HYDROCHLORIDE, OXYMORPHONE HYDROCHLORIDE
 PHENTERMINE HYDROCHLORIDE, PHENTERMINE HYDROCHLORIDE
 TRAZODONE HYDROCHLORIDE, TRAZODONE HYDROCHLORIDE

AUSTARPHARMA

- * AUSTARPHARMA LLC
 FENOFIBRATE (MICRONIZED), FENOFIBRATE
 FENOFIBRATE, FENOFIBRATE

AUSTARPHARMA LLC

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

**** A ****

* AUSTARPHARMA LLC
 METHOCARBAMOL, METHOCARBAMOL
 SERTRALINE HYDROCHLORIDE, SERTRALINE HYDROCHLORIDE

AUXILIUM PHARMS INC

* AUXILIUM PHARMACEUTICALS INC
 TESTOPEL, TESTOSTERONE
 THEO-24, THEOPHYLLINE

AUXILIUM PHARMS LLC

* AUXILIUM PHARMACEUTICALS LLC
 DILATRATE-SR, ISOSORBIDE DINITRATE
 EDEX, ALPROSTADIL
 ROBAXIN-750, METHOCARBAMOL
 SEMPREX-D, ACRIVASTINE
 TESTIM, TESTOSTERONE
 THEO-24, THEOPHYLLINE

AVACOR PRODS

* AVACOR PRODUCTS LLC
 MINOXIDIL EXTRA STRENGTH (FOR MEN), MINOXIDIL (OTC)

AVADEL LEGACY

* AVADEL LEGACY PHARMACEUTICALS LLC
 AKOVAZ, EPHEDRINE SULFATE
 BLOXIVERZ, NEOSTIGMINE METHYLSULFATE
 NOURESS, CYSTEINE HYDROCHLORIDE
 VAZCULEP, PHENYLEPHRINE HYDROCHLORIDE

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

AVEDRO INC

* AVEDRO INC
 PHOTREXA VISCOUS IN DEXTRAN 20%, RIBOFLAVIN 5'-PHOSPHATE SODIUM
 PHOTREXA, RIBOFLAVIN 5'-PHOSPHATE SODIUM

AVEMA PHARMA

* AVEMA PHARMA SOLUTIONS
 IBUPROFEN, IBUPROFEN (OTC)

AVENT

* AVENT INC
 PYTEST KIT, UREA, C-14
 PYTEST, UREA, C-14

AVERITAS

* AVERITAS PHARMA INC
 QUTENZA, CAPSAICIN

AVET

* AVET PHARMACEUTICALS INC
 ACHROMYCIN V, TETRACYCLINE HYDROCHLORIDE

AVEVA

* AVEVA DRUG DELIVERY SYSTEMS INC
 CLONIDINE, CLONIDINE
 FENTANYL-100, FENTANYL
 FENTANYL-12, FENTANYL
 FENTANYL-25, FENTANYL
 FENTANYL-37, FENTANYL
 FENTANYL-50, FENTANYL
 FENTANYL-62, FENTANYL
 FENTANYL-75, FENTANYL

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

* AVEVA DRUG DELIVERY SYSTEMS INC
FENTANYL-87, FENTANYL
NICOTINE, NICOTINE (OTC)

AVID RADIOPHARMS INC

* AVID RADIOPHARMACEUTICALS INC
AMYVID, FLORBETAPIR F-18

AVION PHARMS

* AVION PHARMACEUTICALS LLC
BALCOLTRA, ETHINYL ESTRADIOL
GLOPERBA, COLCHICINE

AVONDALE PHARMS

* AVONDALE PHARMACEUTICALS LLC
NIACOR, NIACIN

AYTU

* AYTU BIOSCIENCE INC
ACIPHEX SPRINKLE, RABEPRAZOLE SODIUM
KARBINAL ER, CARBINOXAMINE MALEATE
TUZISTRA XR, CHLORPHENIRAMINE POLISTIREX
ZOLPIMIST, ZOLPIDEM TARTRATE

**** 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
CEFOTETAN AND DEXTROSE IN DUPLEX CONTAINER, CEFOTETAN DISODIUM
CEFOXITIN AND DEXTROSE IN DUPLEX CONTAINER, CEFOXITIN SODIUM
CEFTAZIDIME IN DEXTROSE CONTAINER, CEFTAZIDIME
CEFTRIAXONE AND DEXTROSE IN DUPLEX CONTAINER, CEFTRIAXONE SODIUM
CEFUROXIME AND DEXTROSE IN DUPLEX CONTAINER, CEFUROXIME 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
DOPAMINE HYDROCHLORIDE AND DEXTROSE 5% IN PLASTIC CONTAINER, DOPAMINE HYDROCHLORIDE
DOPAMINE HYDROCHLORIDE AND DEXTROSE 5%, DOPAMINE HYDROCHLORIDE
FREAMINE HBC 6.9%, AMINO ACIDS
FREAMINE III 10%, AMINO ACIDS
FREAMINE III 3% W/ ELECTROLYTES, AMINO ACIDS
FREAMINE III 8.5% W/ ELECTROLYTES, AMINO ACIDS
FREAMINE III 8.5%, AMINO ACIDS
GLYCINE 1.5% IN PLASTIC CONTAINER, GLYCINE
HEPARIN SODIUM 1,000 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

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** B **

* B BRAUN MEDICAL INC

HEPATAMINE 8%, AMINO ACIDS
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
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
NEPHRAMINE 5.4%, AMINO ACIDS
NUTRILIPID 10%, SOYBEAN OIL
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,

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

**** B ****

*** B BRAUN MEDICAL INC**

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, POTASSIUM CHLORIDE
 PROCALAMINE, AMINO ACIDS
 RESECTISOL IN PLASTIC CONTAINER, MANNITOL
 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
 THEOPHYLLINE 0.04% AND DEXTROSE 5% IN PLASTIC CONTAINER, THEOPHYLLINE
 THEOPHYLLINE 0.08% AND DEXTROSE 5% IN PLASTIC CONTAINER, THEOPHYLLINE
 THEOPHYLLINE 0.16% AND DEXTROSE 5% IN PLASTIC CONTAINER, THEOPHYLLINE
 THEOPHYLLINE 0.32% AND DEXTROSE 5% IN PLASTIC CONTAINER, THEOPHYLLINE
 TROPHAMINE 10%, AMINO ACIDS
 TROPHAMINE, AMINO ACIDS

B BRAUN MEDICAL INC

*** B BRAUN MEDICAL INC**

BUPIVACAINE HYDROCHLORIDE, BUPIVACAINE HYDROCHLORIDE
 CLOROTEKAL, CHLOROPROCAINE HYDROCHLORIDE
 HEPARIN SODIUM, HEPARIN SODIUM
 LIDOCAINE HYDROCHLORIDE, LIDOCAINE HYDROCHLORIDE
 MEROPENEM AND SODIUM CHLORIDE IN DUPLEX CONTAINER, MEROPENEM

BAJAJ

*** BAJAJ MEDICAL LLC**

CHLORHEXIDINE GLUCONATE, CHLORHEXIDINE GLUCONATE (OTC)

BARR

*** BARR LABORATORIES INC**

AMILORIDE HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, AMILORIDE HYDROCHLORIDE
 ANAGRELIDE HYDROCHLORIDE, ANAGRELIDE HYDROCHLORIDE
 ARANELLE, ETHINYL ESTRADIOL
 ASPIRIN AND DIPYRIDAMOLE, ASPIRIN
 BALZIVA-28, ETHINYL ESTRADIOL
 CHLORDIAZEPOXIDE HYDROCHLORIDE, CHLORDIAZEPOXIDE HYDROCHLORIDE
 CLONAZEPAM, CLONAZEPAM
 DANAZOL, DANAZOL
 DEMECLOCYCLINE HYDROCHLORIDE, DEMECLOCYCLINE HYDROCHLORIDE
 DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE
 DEXTROAMPHETAMINE SULFATE, DEXTROAMPHETAMINE SULFATE
 DIAZEPAM, DIAZEPAM
 DIPYRIDAMOLE, DIPYRIDAMOLE
 DROSPIRENONE AND ETHINYL ESTRADIOL, DROSPIRENONE
 DUTASTERIDE, DUTASTERIDE
 ESTRADIOL AND NORETHINDRONE ACETATE, ESTRADIOL
 ESTRADIOL AND NORGESTIMATE, ESTRADIOL
 ETHAMBUTOL HYDROCHLORIDE, ETHAMBUTOL HYDROCHLORIDE
 FEXOFENADINE HYDROCHLORIDE, FEXOFENADINE HYDROCHLORIDE
 FLUDROCORTISONE ACETATE, FLUDROCORTISONE ACETATE
 FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 GALANTAMINE HYDROBROMIDE, GALANTAMINE HYDROBROMIDE
 HYDROMORPHONE HYDROCHLORIDE, HYDROMORPHONE HYDROCHLORIDE
 HYDROXYUREA, HYDROXYUREA
 HYDROXYZINE PAMOATE, HYDROXYZINE PAMOATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

**** B ****

- * BARR LABORATORIES INC
 - ISONIAZID, ISONIAZID
 - JUNEL 1.5/30, ETHINYL ESTRADIOL
 - JUNEL 1/20, ETHINYL ESTRADIOL
 - JUNEL FE 1.5/30, ETHINYL ESTRADIOL
 - JUNEL FE 1/20, ETHINYL ESTRADIOL
 - KARIVA, DESOGESTREL
 - KELNOR, ETHINYL ESTRADIOL
 - LEFLUNOMIDE, LEFLUNOMIDE
 - LESSINA-28, ETHINYL ESTRADIOL
 - MEDROXYPROGESTERONE ACETATE, MEDROXYPROGESTERONE ACETATE
 - MEFLOQUINE HYDROCHLORIDE, MEFLOQUINE HYDROCHLORIDE
 - MEGESTROL ACETATE, MEGESTROL ACETATE
 - METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 - 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
 - ONDANSETRON, ONDANSETRON
 - PHENTERMINE HYDROCHLORIDE, PHENTERMINE HYDROCHLORIDE
 - PORTIA-28, ETHINYL ESTRADIOL
 - SPRINTEC, ETHINYL ESTRADIOL
 - TEMOZOLOMIDE, TEMOZOLOMIDE
 - TREXALL, METHOTREXATE SODIUM
 - TRI-LEGEST 21, ETHINYL ESTRADIOL
 - TRI-LEGEST FE, ETHINYL ESTRADIOL
 - TRI-SPRINTEC, ETHINYL ESTRADIOL
 - WARFARIN SODIUM, WARFARIN SODIUM

- * BARR PHARMACEUTICALS
 - LEUCOVORIN CALCIUM, LEUCOVORIN CALCIUM

BARR LABS DIV TEVA

- * BARR LABORATORIES INC SUB TEVA PHARMACEUTICALS USA
 - BUDESONIDE, BUDESONIDE

BARR LABS INC

- * BARR LABORATORIES INC
 - ACITRETIN, ACITRETIN
 - CLOZAPINE, CLOZAPINE
 - DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE
 - ESTRADIOL, ESTRADIOL
 - METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
 - MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE
 - NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL AND FERROUS FUMARATE, ETHINYL ESTRADIOL
 - NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
 - OLANZAPINE, OLANZAPINE
 - OLOPATADINE HYDROCHLORIDE, OLOPATADINE HYDROCHLORIDE
 - OXYCODONE HYDROCHLORIDE AND IBUPROFEN, IBUPROFEN
 - TRETINOIN, TRETINOIN
 - TRI LO SPRINTEC, ETHINYL ESTRADIOL

BAUDAX

- * BAUDAX BIO INC
 - ANJESO, MELOXICAM

BAUSCH

- * BAUSCH HEALTH AMERICAS INC
 - ACANYA, BENZOYL PEROXIDE
 - ARAZLO, TAZAROTENE
 - BENZAFLIN, BENZOYL PEROXIDE
 - BRYHALI, HALOBETASOL PROPIONATE
 - DUOBRII, HALOBETASOL PROPIONATE
 - EDECIN, ETHACRYNATE SODIUM
 - EDECIN, ETHACRYNIC ACID

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** B ****

- * BAUSCH HEALTH AMERICAS INC
 - EFUDEX, FLUOROURACIL
 - JUBLIA, EFINAACONAZOLE
 - LOCOID, HYDROCORTISONE BUTYRATE
 - MEPHYTON, PHYTONADIONE
 - ONEXTON, BENZOYL PEROXIDE
 - OXSORALEN-ULTRA, METHOXSALEN
 - SYPRINE, TRIENTINE HYDROCHLORIDE
- * BAUSCH HEALTH IRELAND LTD
 - ERTACZO, SERTACONAZOLE NITRATE
 - TARGRETIN, BEXAROTENE
 - TETRACAINE HYDROCHLORIDE, TETRACAINE HYDROCHLORIDE
- * BAUSCH HEALTH US LLC
 - ALDARA, IMIQUIMOD
 - ANCOBON, FLUCYTOSINE
 - ATIVAN, LORAZEPAM
 - CARDIZEM CD, DILTIAZEM HYDROCHLORIDE
 - CARDIZEM LA, DILTIAZEM HYDROCHLORIDE
 - CARDIZEM, DILTIAZEM HYDROCHLORIDE
 - CLINDAGEL, CLINDAMYCIN PHOSPHATE
 - D.H.E. 45, DIHYDROERGOTAMINE MESYLATE
 - DEMSEER, METYROSINE
 - DIASTAT ACUDIAL, DIAZEPAM
 - DIASTAT, DIAZEPAM
 - ELIDEL, PIMECROLIMUS
 - ISORDIL, ISOSORBIDE DINITRATE
 - ISUPREL, ISOPROTERENOL HYDROCHLORIDE
 - LOCOID, HYDROCORTISONE BUTYRATE
 - LOPROX, CICLOPIROX
 - MESTINON, PYRIDOSTIGMINE BROMIDE
 - METROGEL-VAGINAL, METRONIDAZOLE
 - MIGRANAL, DIHYDROERGOTAMINE MESYLATE
 - MINOCIN, MINOCYCLINE HYDROCHLORIDE
 - TASMAR, TOLCAPONE
 - TIAZAC, DILTIAZEM HYDROCHLORIDE
 - VASERETIC, ENALAPRIL MALEATE
 - VASOTEC, ENALAPRIL MALEATE
 - XERESE, ACYCLOVIR
 - ZELAPAR, SELEGILINE HYDROCHLORIDE
 - ZOVIRAX, ACYCLOVIR
 - ZYCLARA, IMIQUIMOD

BAUSCH AND LOMB

- * BAUSCH AND LOMB INC
 - ALAWAY, KETOTIFEN FUMARATE (OTC)
 - ALREX, LOTEHPREDNOL ETABONATE
 - BESIVANCE, BESIFLOXACIN HYDROCHLORIDE
 - DICLOFENAC SODIUM, DICLOFENAC SODIUM
 - DORZOLAMIDE HYDROCHLORIDE AND TIMOLOL MALEATE, DORZOLAMIDE HYDROCHLORIDE
 - DORZOLAMIDE HYDROCHLORIDE, DORZOLAMIDE HYDROCHLORIDE
 - FLURBIPROFEN SODIUM, FLURBIPROFEN SODIUM
 - IPRATROPIUM BROMIDE, IPRATROPIUM BROMIDE
 - ISTALOL, TIMOLOL MALEATE
 - LATANOPROST, LATANOPROST
 - LOTEMAX, LOTEHPREDNOL 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
 - TIMOLOL MALEATE, TIMOLOL MALEATE
 - TROPICAMIDE, TROPICAMIDE
 - VITRASE, HYALURONIDASE
 - VYZULTA, LATANOPROSTENE BUNOD
 - ZIRGAN, GANCICLOVIR

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** B ****

- * BAUSCH AND LOMB INC
ZYLET, LOTEPREDNOL ETABONATE
- * BAUSCH AND LOMB PHARMACEUTICALS INC
BACITRACIN ZINC AND POLYMYXIN B SULFATE, BACITRACIN ZINC
BRIMONIDINE TARTRATE, BRIMONIDINE TARTRATE
CROMOLYN SODIUM, CROMOLYN SODIUM (OTC)
DESMOPRESSIN ACETATE, DESMOPRESSIN ACETATE
DEXAMETHASONE SODIUM PHOSPHATE, DEXAMETHASONE SODIUM PHOSPHATE
DEXASPORIN, DEXAMETHASONE
ERYTHROMYCIN, ERYTHROMYCIN
FLUNISOLIDE, FLUNISOLIDE
GENTAMICIN SULFATE, GENTAMICIN SULFATE
IPRATROPIUM BROMIDE, IPRATROPIUM BROMIDE
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
TIMOLOL MALEATE, TIMOLOL MALEATE
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
BEPREVE, BEPOTASTINE BESILATE
LOTEMAX SM, LOTEPREDNOL ETABONATE
LOTEMAX, LOTEPREDNOL ETABONATE
LUMIFY, BRIMONIDINE TARTRATE (OTC)

BAXTER HLTHCARE

- * BAXTER HEALTHCARE CORP
ACETIC ACID 0.25% IN PLASTIC CONTAINER, ACETIC ACID, GLACIAL
AMINOACETIC ACID 1.5% IN PLASTIC CONTAINER, GLYCINE
ANCEF IN PLASTIC CONTAINER, CEFAZOLIN SODIUM
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 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,
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

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

**** B ****

* BAXTER HEALTHCARE CORP

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
 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
 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

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** B **

- * BAXTER HEALTHCARE CORP
 MESNEX, MESNA
 MILRINONE LACTATE IN DEXTROSE 5% IN PLASTIC CONTAINER, MILRINONE LACTATE
 NALLPEN IN PLASTIC CONTAINER, NAFCILLIN 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% IN PLASTIC CONTAINER, POTASSIUM
 POTASSIUM CHLORIDE 0.15% IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, POTASSIUM
 POTASSIUM CHLORIDE 0.3% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, POTASSIUM
 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
 VANCOCIN HYDROCHLORIDE IN PLASTIC CONTAINER, VANCOMYCIN HYDROCHLORIDE
- * BAXTER HEALTHCARE INTERNATIONAL SPECIALTY THERAPIES DIV
 PROSOL 20% SULFITE FREE IN PLASTIC CONTAINER, AMINO ACIDS
- BAXTER HLTHCARE CORP**
- * BAXTER HEALTHCARE CORP
 BIVALIRUDIN IN 0.9% SODIUM CHLORIDE, BIVALIRUDIN

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** B ****

- * BAXTER HEALTHCARE CORP
 BUPIVACAINE HYDROCHLORIDE, BUPIVACAINE HYDROCHLORIDE
 CEFAZOLIN IN PLASTIC CONTAINER, CEFAZOLIN SODIUM
 CIPROFLOXACIN IN DEXTROSE 5% IN PLASTIC CONTAINER, CIPROFLOXACIN
 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
 DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE
 DOXIL (LIPOSOMAL), DOXORUBICIN HYDROCHLORIDE
 EPTIFIBATIDE, EPTIFIBATIDE
 FLUCONAZOLE IN SODIUM CHLORIDE 0.9%, FLUCONAZOLE
 FOSAPREPITANT DIMEGLUMINE, FOSAPREPITANT DIMEGLUMINE
 FUROSEMIDE, FUROSEMIDE
 KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
 LEVOFLOXACIN IN DEXTROSE 5% IN PLASTIC CONTAINER, LEVOFLOXACIN
 LEVOFLOXACIN, LEVOFLOXACIN
 METOPROLOL TARTRATE, METOPROLOL TARTRATE
 METRONIDAZOLE IN PLASTIC CONTAINER, METRONIDAZOLE
 MYXREDLIN, INSULIN HUMAN
 NOREPINEPHRINE BITARTRATE, NOREPINEPHRINE BITARTRATE
 ONDANSETRON HYDROCHLORIDE PRESERVATIVE FREE, ONDANSETRON HYDROCHLORIDE
 ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
 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
 TOBRAMYCIN SULFATE, TOBRAMYCIN SULFATE
- * BAXTER HEALTHCARE CORP ANESTHESIA AND CRITICAL CARE
 PROTOPAM CHLORIDE, PRALIDOXIME CHLORIDE

BAYER

- * BAYER HEALTHCARE LLC
 ALEVE, NAPROXEN SODIUM (OTC)
 ALEVE-D SINUS & COLD, NAPROXEN SODIUM (OTC)

BAYER HEALTHCARE

- * BAYER HEALTHCARE PHARMACEUTICALS INC
 ALIQOPA, COPANLISIB DIHYDROCHLORIDE
 NUBEQA, DAROLUTAMIDE
 VITRAKVI, LAROTRECTINIB SULFATE

BAYER HEALTHCARE LLC

- * BAYER HEALTHCARE LLC
 CHILDREN'S CLARITIN, LORATADINE (OTC)
 CLARITIN HIVES RELIEF REDITAB, LORATADINE (OTC)
 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)
 ZEGERID OTC, OMEPRAZOLE (OTC)

BAYER HLTHCARE

- * BAYER HEALTHCARE CONSUMER CARE
 ALEVE PM, DIPHENHYDRAMINE HYDROCHLORIDE (OTC)
- * BAYER HEALTHCARE PHARMACEUTICALS INC
 ADEMPAS, RIOCIGUAT
 ANGELIQ, DROSPIRENONE
 AVELOX, MOXIFLOXACIN HYDROCHLORIDE
 BEYAZ, DROSPIRENONE
 BILTRICIDE, PRAZIQUANTEL

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

**** B ****

* BAYER HEALTHCARE PHARMACEUTICALS INC
 CIPRO, CIPROFLOXACIN
 CIPRO, CIPROFLOXACIN HYDROCHLORIDE
 CLIMARA PRO, ESTRADIOL
 CLIMARA, ESTRADIOL
 EOVI, GADOXETATE DISODIUM
 GADAVIST, GADOBUTROL
 KYLEENA, LEVONORGESTREL
 LEVITRA, VARDENAFIL HYDROCHLORIDE
 MENOSTAR, ESTRADIOL
 MIRENA, LEVONORGESTREL
 NATAZIA, DIENOGEST
 NEXAVAR, SORAFENIB TOSYLATE
 SAFYRAL, DROSPIRENONE
 SKYLA, LEVONORGESTREL
 STAXYN, VARDENAFIL HYDROCHLORIDE
 STIVARGA, REGORAFENIB
 ULTRAVIST (PHARMACY BULK), IOPROMIDE
 ULTRAVIST 300, IOPROMIDE
 ULTRAVIST 370, IOPROMIDE
 VITRAKVI, LAROTRECTINIB SULFATE
 XOFIGO, RADIUM RA-223 DICHLORIDE
 YASMIN, DROSPIRENONE
 YAZ, DROSPIRENONE

BAYSHORE PHARMS LLC

* BAYSHORE PHARMACEUTICALS LLC
 DIPHENOXYLATE HYDROCHLORIDE AND ATROPINE SULFATE, ATROPINE SULFATE
 METHSCOPOLAMINE BROMIDE, METHSCOPOLAMINE BROMIDE
 PINDOLOL, PINDOLOL
 PRIMAQUINE PHOSPHATE, PRIMAQUINE PHOSPHATE

BDSI

* BIODELIVERY SCIENCES INTERNATIONAL INC
 BELBUCA, BUPRENORPHINE HYDROCHLORIDE
 BUNAVAIL, BUPRENORPHINE HYDROCHLORIDE
 SYMPROIC, NALDEMEDINE TOSYLATE

BE PHARMS

* BE PHARMACEUTICALS AG
 DAPTOMYCIN, DAPTOMYCIN
 FOSAPREPITANT DIMEGLUMINE, FOSAPREPITANT DIMEGLUMINE
 NEOSTIGMINE METHYLSULFATE, NEOSTIGMINE METHYLSULFATE

BECTON DICKINSON

* BECTON DICKINSON AND CO
 CHLORHEXIDINE GLUCONATE, CHLORHEXIDINE GLUCONATE (OTC)
 E-Z SCRUB 201, POVIDONE-IODINE (OTC)
 E-Z SCRUB 241, POVIDONE-IODINE (OTC)

BECTON DICKINSON CO

* BECTON DICKINSON AND CO
 CHLORAPREP ONE-STEP FREPP, CHLORHEXIDINE GLUCONATE (OTC)
 CHLORAPREP ONE-STEP SEPP, CHLORHEXIDINE GLUCONATE (OTC)
 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 YILING

* BEIJING YILING BIO-ENGINEERING AND TECHNOLOGY CO LTD
 ANASTROZOLE, ANASTROZOLE
 LETROZOLE, LETROZOLE

BELCHER

* BELCHER PHARMACEUTICALS LLC
 ABLYSINOL, ALCOHOL
 CEFIXIME, CEFIXIME
 EPINEPHRINE, EPINEPHRINE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

**** B ****

* BELCHER PHARMACEUTICALS LLC
 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 INC

* BELOTECA INC
 DIAZEPAM, DIAZEPAM

BENUVIA

* BENUVIA THERAPEUTICS INC
 SYNDROS, DRONABINOL

BEXIMCO PHARMS USA

* BEXIMCO PHARMACEUTICALS USA INC
 CYPROHEPTADINE HYDROCHLORIDE, CYPROHEPTADINE HYDROCHLORIDE
 LOVASTATIN, LOVASTATIN
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 METHOCARBAMOL, METHOCARBAMOL
 NADOLOL, NADOLOL
 SOTALOL HYDROCHLORIDE, SOTALOL HYDROCHLORIDE

BEXIMCO USA

* BEXIMCO PHARMACEUTICALS USA INC
 CARVEDILOL, CARVEDILOL

BIOCODEX SA

* BIOCODEX SA
 DIACOMIT, STIRIPENTOL

BIOCON LIMITED

* BIOCON LIMITED
 SIMVASTATIN, SIMVASTATIN

BIOCON LTD

* BIOCON LTD
 FINGOLIMOD HYDROCHLORIDE, FINGOLIMOD HYDROCHLORIDE
 ROSUVASTATIN CALCIUM, ROSUVASTATIN CALCIUM

BIOCRIST

* BIOCRIST PHARMACEUTICALS INC
 RAPIVAB, PERAMIVIR

BIOFRONTERA

* BIOFRONTERA BIOSCIENCE GMBH
 AMELUZ, AMINOLEVULINIC ACID HYDROCHLORIDE

BIOGEN

* BIOGEN
 VUMERITY, DIROXIMEL FUMARATE

BIOGEN IDEC

* BIOGEN IDEC INC
 SPINRAZA, NUSINERSEN SODIUM

BIOGEN IDEC INC

* BIOGEN IDEC INC
 TECFIDERA, DIMETHYL FUMARATE

BIOHAVEN PHARM

* BIOHAVEN PHARMACEUTICAL HOLDING CO LTD
 NURTEC ODT, RIMEGEPANT SULFATE

BIOMARIN PHARM

* BIOMARIN PHARMACEUTICAL INC
 KUVAN, SAPROPTERIN DIHYDROCHLORIDE

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 INC

* BIONPHARMA INC

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

**** B ****

* BIONPHARMA INC
 AMANTADINE HYDROCHLORIDE, AMANTADINE HYDROCHLORIDE
 AMPHETAMINE SULFATE, AMPHETAMINE SULFATE
 AZITHROMYCIN, AZITHROMYCIN
 BENZONATATE, BENZONATATE
 BEXAROTENE, BEXAROTENE
 CALCITRIOL, CALCITRIOL
 CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 CETIRIZINE HYDROCHLORIDE HIVES RELIEF, CETIRIZINE HYDROCHLORIDE (OTC)
 CLOBAZAM, CLOBAZAM
 COLESEVELAM HYDROCHLORIDE, COLESEVELAM HYDROCHLORIDE
 DEFERASIROX, DEFERASIROX
 DICLOFENAC POTASSIUM, DICLOFENAC POTASSIUM
 DOFETILIDE, DOFETILIDE
 DUTASTERIDE, DUTASTERIDE
 ETHOSUXIMIDE, ETHOSUXIMIDE
 GRANISETRON HYDROCHLORIDE PRESERVATIVE FREE, GRANISETRON HYDROCHLORIDE
 GRANISETRON HYDROCHLORIDE, GRANISETRON HYDROCHLORIDE
 IBUPROFEN AND DIPHENHYDRAMINE HYDROCHLORIDE, DIPHENHYDRAMINE HYDROCHLORIDE (OTC)
 IBUPROFEN, IBUPROFEN (OTC)
 LOPERAMIDE HYDROCHLORIDE, LOPERAMIDE HYDROCHLORIDE (OTC)
 METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
 MIDOL LIQUID GELS, IBUPROFEN (OTC)
 NAPROXEN SODIUM, NAPROXEN SODIUM (OTC)
 NIMODIPINE, NIMODIPINE
 OMEGA-3-ACID ETHYL ESTERS, OMEGA-3-ACID ETHYL ESTERS
 OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE
 PARICALCITOL, PARICALCITOL
 PROGESTERONE, PROGESTERONE
 RUFINAMIDE, RUFINAMIDE
 TETRABENAZINE, TETRABENAZINE
 VALPROIC ACID, VALPROIC ACID
 VITAMIN D, ERGOCALCIFEROL
 ZONISAMIDE, ZONISAMIDE

BLAIREX

* BLAIREX LABORATORIES INC
 BRONCHO SALINE, SODIUM CHLORIDE (OTC)

BLUE EARTH

* BLUE EARTH DIAGNOSTICS LTD
 AXUMIN, FLUCICLOVINE F-18

BLUEPRINT MEDICINES

* BLUEPRINT MEDICINES CORP
 AYVAKIT, AVAPRITINIB

BOEHRINGER INGELHEIM

* BOEHRINGER INGELHEIM
 CATAPRES, CLONIDINE HYDROCHLORIDE
 CATAPRES-TTS-1, CLONIDINE
 CATAPRES-TTS-2, CLONIDINE
 CATAPRES-TTS-3, CLONIDINE
 GILOTRIF, AFATINIB DIMALEATE
 GLYXAMBI, EMPAGLIFLOZIN
 MICARDIS HCT, HYDROCHLOROTHIAZIDE
 MICARDIS, TELMISARTAN
 MIRAPEX, PRAMIPEXOLE DIHYDROCHLORIDE
 * BOEHRINGER INGELHEIM PHARMACEUTICALS INC
 AGGRENOX, ASPIRIN
 APTIVUS, TIPRANAVIR
 ATROVENT HFA, IPRATROPIUM BROMIDE
 COMBIVENT RESPIMAT, ALBUTEROL SULFATE
 JARDIANCE, EMPAGLIFLOZIN
 JENTADUETO XR, LINAGLIPTIN
 JENTADUETO, LINAGLIPTIN
 MIRAPEX ER, PRAMIPEXOLE DIHYDROCHLORIDE
 MOBIC, MELOXICAM
 OFEV, NINTEDANIB ESYLATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

**** B ****

* BOEHRINGER INGELHEIM PHARMACEUTICALS INC
 PERSANTINE, DIPYRIDAMOLE
 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
 TWYNSTA, AMLODIPINE BESYLATE
 VIRAMUNE XR, NEVIRAPINE
 VIRAMUNE, NEVIRAPINE

BOSCOGEN

* BOSCOGEN INC
 ARIPIPRAZOLE, ARIPIPRAZOLE
 CAPTOPRIL, CAPTOPRIL
 CYPROHEPTADINE HYDROCHLORIDE, CYPROHEPTADINE HYDROCHLORIDE
 REPAGLINIDE, REPAGLINIDE
 SELEGILINE HYDROCHLORIDE, SELEGILINE HYDROCHLORIDE

BPI LABS LLC

* BPI LABS LLC
 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
 GASTROGRAFIN, DIATRIZOATE MEGLUMINE
 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

BRAINTREE

* BRAINTREE LABORATORIES INC
 GOLYTELY, POLYETHYLENE GLYCOL 3350
 NULYTELY, POLYETHYLENE GLYCOL 3350
 NULYTELY-FLAVORED, POLYETHYLENE GLYCOL 3350

BRAINTREE LABS

* BRAINTREE LABORATORIES INC
 PIZENSY, LACTITOL
 SUPREP BOWEL PREP KIT, MAGNESIUM SULFATE

BRECKENRIDGE

* BRECKENRIDGE PHARMACEUTICAL INC

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

**** B ****

* BRECKENRIDGE PHARMACEUTICAL INC
 ALPRAZOLAM, ALPRAZOLAM
 AZELASTINE HYDROCHLORIDE, AZELASTINE HYDROCHLORIDE
 BRETILUM TOSYLATE, BRETILUM TOSYLATE
 CETIRIZINE HYDROCHLORIDE, CETIRIZINE HYDROCHLORIDE
 CLOBAZAM, CLOBAZAM
 DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE
 ENTECAVIR, ENTECAVIR
 EPINASTINE HYDROCHLORIDE, EPINASTINE HYDROCHLORIDE
 EPLERENONE, EPLERENONE
 EXEMESTANE, EXEMESTANE
 IMATINIB MESYLATE, IMATINIB MESYLATE
 LEVETIRACETAM, LEVETIRACETAM
 MEGESTROL ACETATE, MEGESTROL ACETATE
 METHYLERGONOVINE MALEATE, METHYLERGONOVINE MALEATE
 METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
 NEOMYCIN SULFATE, NEOMYCIN SULFATE
 OMEPRAZOLE, OMEPRAZOLE
 PEG-3350, POTASSIUM CHLORIDE, SODIUM BICARBONATE, SODIUM CHLORIDE, POLYETHYLENE GLYCOL
 PIOGLITAZONE HYDROCHLORIDE, PIOGLITAZONE HYDROCHLORIDE
 RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
 RIVASTIGMINE, RIVASTIGMINE
 ROFLUMILAST, ROFLUMILAST
 SOLIFENACIN SUCCINATE, SOLIFENACIN SUCCINATE
 SUCCINYLCHOLINE CHLORIDE, SUCCINYLCHOLINE CHLORIDE
 TETRACYCLINE HYDROCHLORIDE, TETRACYCLINE HYDROCHLORIDE
 ZOLEDRONIC ACID, ZOLEDRONIC ACID

BRECKENRIDGE PHARM

* BRECKENRIDGE PHARMACEUTICAL INC
 CILOSTAZOL, CILOSTAZOL
 ESTRADIOL AND NORETHINDRONE ACETATE, ESTRADIOL
 LEVETIRACETAM, LEVETIRACETAM
 MELOXICAM, MELOXICAM
 OXCARBAZEPINE, OXCARBAZEPINE
 PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE
 TERBINAFINE HYDROCHLORIDE, TERBINAFINE 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

BRISTOL MYERS SQUIBB

* BRISTOL MYERS SQUIBB
 AZACTAM, AZTREONAM
 BARACLUDE, ENTECAVIR
 PRAVACHOL, PRAVASTATIN SODIUM

* BRISTOL MYERS SQUIBB CO
 DROXIA, HYDROXYUREA
 HYDREA, HYDROXYUREA
 REYATAZ, ATAZANAVIR SULFATE
 SPRYCEL, DASATINIB
 SUSTIVA, EFAVIRENZ
 VIDEX EC, DIDANOSINE

* BRISTOL MYERS SQUIBB CO PHARMACEUTICAL RESEARCH INSTITUTE
 ELIQUIS, APIXABAN
 ZERIT, STAVUDINE

* BRISTOL MYERS SQUIBB PHARMA CO
 COUMADIN, WARFARIN SODIUM

BRISTOL-MYERS SQUIBB

* BRISTOL-MYERS SQUIBB CO
 EVOTAZ, ATAZANAVIR SULFATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** B ****

* BRISTOL-MYERS SQUIBB CO
VIDEX, DIDANOSINE

BTCP PHARMA

* BTCP PHARMA LLC
LAZANDA, FENTANYL CITRATE
SUBSYS, FENTANYL

BWXT ITG

* BWXT ITG CANADA INC
INDIUM IN 111 OXYQUINOLINE, INDIUM IN-111 OXYQUINOLINE

**** C ******CADILA**

* CADILA HEALTHCARE LTD
ACETAZOLAMIDE, ACETAZOLAMIDE
ACYCLOVIR, ACYCLOVIR
CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
DESONIDE, DESONIDE
DICLOFENAC SODIUM, DICLOFENAC SODIUM
DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
DUTASTERIDE, DUTASTERIDE
ESCITALOPRAM OXALATE, ESCITALOPRAM OXALATE
FELBAMATE, FELBAMATE
FLUOCINONIDE, FLUOCINONIDE
GEMFIBROZIL, GEMFIBROZIL
INDOMETHACIN, INDOMETHACIN
MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
METRONIDAZOLE, METRONIDAZOLE
MODAFINIL, MODAFINIL
NYSTATIN, NYSTATIN
PIROXICAM, PIROXICAM
RANOLAZINE, RANOLAZINE
ROPINIROLE HYDROCHLORIDE, ROPINIROLE 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
FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
FOLIC ACID, FOLIC ACID
GEMFIBROZIL, GEMFIBROZIL
GLYBURIDE, GLYBURIDE
HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
LABETALOL HYDROCHLORIDE, LABETALOL HYDROCHLORIDE
METRONIDAZOLE, METRONIDAZOLE
NATEGLINIDE, NATEGLINIDE
OFLOXACIN, OFLOXACIN
RIVASTIGMINE TARTRATE, RIVASTIGMINE TARTRATE
ROSUVASTATIN CALCIUM, ROSUVASTATIN CALCIUM
TELMISARTAN, TELMISARTAN
TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE
VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE
VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE

CADISTA PHARMS

* CADISTA PHARMACEUTICALS INC
LOSARTAN POTASSIUM AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
LOSARTAN POTASSIUM, LOSARTAN POTASSIUM

CALL INC

* CALL INC DBA ROCHESTER PHARMACEUTICALS
ADAPALENE, ADAPALENE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** C ******CAPELLON PHARMS LLC**

* CAPELLON PHARMACEUTICALS LLC
POLMON, DEXCHLORPHENIRAMINE MALEATE

CAPLIN

* CAPLIN STERILES LTD
GLYCOPYRROLATE, GLYCOPYRROLATE
SODIUM NITROPRUSSIDE, SODIUM NITROPRUSSIDE

CARDINAL HEALTH 414

* CARDINAL HEALTH 414 LLC CARDINAL HEALTH NUCLEAR PHARMACY SERVICES
AMMONIA N 13, AMMONIA N-13
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

CARDINAL HEALTH 418

* CARDINAL HEALTH 418 INC
SODIUM IODIDE I 123, SODIUM IODIDE I-123

CARIBE HOLDINGS

* CARIBE HOLDINGS CAYMAN CO LTD DBA PURACAP CARIBE
DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
GEMFIBROZIL, GEMFIBROZIL

CARLSBAD

* CARLSBAD TECHNOLOGY INC
ACYCLOVIR, ACYCLOVIR
CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE
DICLOFENAC SODIUM, DICLOFENAC SODIUM
FAMOTIDINE, FAMOTIDINE
GLIMEPIRIDE, GLIMEPIRIDE
LOVASTATIN, LOVASTATIN

CARLSBAD TECHNOLOGY

* CARLSBAD TECHNOLOGY INC
ACYCLOVIR, ACYCLOVIR

CASI PHARMS INC

* CASI PHARMACEUTICALS INC
BENAZEPRIL HYDROCHLORIDE, BENAZEPRIL HYDROCHLORIDE
BISOPROLOL FUMARATE, BISOPROLOL FUMARATE
BUPRENORPHINE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE
CEFPROZIL, CEFPROZIL
CILOSTAZOL, CILOSTAZOL
DESVENLAFAXINE SUCCINATE, DESVENLAFAXINE SUCCINATE
DICLOFENAC POTASSIUM, DICLOFENAC POTASSIUM
ENTECAVIR, ENTECAVIR
HEPARIN SODIUM, HEPARIN SODIUM
LISINAPRIL, LISINAPRIL
METHIMAZOLE, METHIMAZOLE
NABUMETONE, NABUMETONE
ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
REPAGLINIDE, REPAGLINIDE
TENOFIVIR DISOPROXIL FUMARATE, TENOFIVIR DISOPROXIL FUMARATE
TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE

CASPER PHARMA LLC

* CASPER PHARMA LLC
ANTIVERT, MECLIZINE HYDROCHLORIDE
AQUASOL A, VITAMIN A PALMITATE
CASPORYN HC, HYDROCORTISONE
FURADANTIN, NITROFURANTOIN
NEOSPORIN, BACITRACIN ZINC
ZYLOPRIM, ALLOPURINOL

CATALENT

* CATALENT PHARMA SOLUTIONS LLC
NAPROXEN SODIUM, NAPROXEN SODIUM (OTC)
VALPROIC ACID, VALPROIC ACID

CATALYST PHARMS

* CATALYST PHARMACEUTICALS INC

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

**** C ****

* CATALYST PHARMACEUTICALS INC
FIRDAPSE, AMIFAMPRIDINE PHOSPHATE

CEDIPROF INC

* CEDIPROF INC
DEXMETHYLPHENIDATE HYDROCHLORIDE, DEXMETHYLPHENIDATE HYDROCHLORIDE
LEVO-T, LEVOTHYROXINE SODIUM **

CELATOR PHARMS

* CELATOR PHARMACEUTICALS INC
VYXEOS, CYTARABINE

CELGENE

* CELGENE CORP
ISTODAX, ROMIDEPSIN
POMALYST, POMALIDOMIDE
REVLIMID, LENALIDOMIDE
THALOMID, THALIDOMIDE
VIDAZA, AZACITIDINE

CELGENE CORP

* CELGENE CORP
IDHIFA, ENASIDENIB MESYLATE

CELLTRION

* CELLTRION INC
CLOPIDOGREL BISULFATE, CLOPIDOGREL BISULFATE
DIVALPROEX SODIUM, DIVALPROEX SODIUM
FAMOTIDINE, FAMOTIDINE
LEVOFLOXACIN, LEVOFLOXACIN
LINEZOLID, LINEZOLID
MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
PIOGLITAZONE HYDROCHLORIDE, PIOGLITAZONE HYDROCHLORIDE
RISPERIDONE, RISPERIDONE
RIZATRIPTAN BENZOATE, RIZATRIPTAN BENZOATE
ROPINIROLE HYDROCHLORIDE, ROPINIROLE HYDROCHLORIDE
SOLIFENACIN SUCCINATE, SOLIFENACIN SUCCINATE
TEMIXYS, LAMIVUDINE
ZONISAMIDE, ZONISAMIDE

CEPHALON

* CEPHALON INC
ACTIQ, FENTANYL CITRATE
FENTORA, FENTANYL CITRATE
GABITRIL, TIAGABINE HYDROCHLORIDE
NUVIGIL, ARMODAFINIL
PROVIGIL, MODAFINIL
TREANDA, BENDAMUSTINE HYDROCHLORIDE
TRISENOX, ARSENIC TRIOXIDE

CEROVENE INC

* CEROVENE INC
AMPHETAMINE SULFATE, AMPHETAMINE SULFATE
PYRIMETHAMINE, PYRIMETHAMINE

CHANGZHOU PHARM

* CHANGZHOU PHARMACEUTICAL FACTORY
DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
ROSUVASTATIN CALCIUM, ROSUVASTATIN CALCIUM

CHARTWELL

* CHARTWELL LIFE MOLECULES LLC
CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE
CLARITHROMYCIN, CLARITHROMYCIN
FLUCONAZOLE, FLUCONAZOLE

CHARTWELL LIFE SCI

* CHARTWELL LIFE SCIENCE LLC
DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
DOXYCYCLINE, DOXYCYCLINE
METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE

CHARTWELL MOLECULAR

* CHARTWELL MOLECULAR HOLDINGS LLC
CALCIUM ACETATE, CALCIUM ACETATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** C ****

* CHARTWELL MOLECULAR HOLDINGS LLC
 CARVEDILOL, CARVEDILOL
 CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE
 FOLIC ACID, FOLIC ACID
 GRANISETRON HYDROCHLORIDE, GRANISETRON HYDROCHLORIDE
 IRBESARTAN, IRBESARTAN
 RAMIPRIL, RAMIPRIL
 RISPERIDONE, RISPERIDONE

CHARTWELL MOLECULES

* CHARTWELL MOLECULES LLC
 DISULFIRAM, DISULFIRAM
 GEMFIBROZIL, GEMFIBROZIL
 NABUMETONE, NABUMETONE
 SULFAMETHOXAZOLE AND TRIMETHOPRIM, SULFAMETHOXAZOLE

CHARTWELL RX

* CHARTWELL RX SCIENCES LLC
 CALCIUM ACETATE, CALCIUM ACETATE
 CILOSTAZOL, CILOSTAZOL
 DUVOID, BETHANECHOL CHLORIDE
 GRISEOFULVIN, GRISEOFULVIN, MICROSIZE
 INDOMETHACIN, INDOMETHACIN
 LEVETIRACETAM, LEVETIRACETAM
 MOEXIPRIL HYDROCHLORIDE, MOEXIPRIL HYDROCHLORIDE
 PRAVASTATIN SODIUM, PRAVASTATIN SODIUM
 PREDNISOLONE SODIUM PHOSPHATE, PREDNISOLONE SODIUM PHOSPHATE
 RIVASTIGMINE TARTRATE, RIVASTIGMINE TARTRATE

CHARTWELL TETRA

* CHARTWELL TETRA LLC
 TETRACYCLINE HYDROCHLORIDE, TETRACYCLINE HYDROCHLORIDE

CHATTEM

* CHATTEM INC
 UNISOM, DOXYLAMINE SUCCINATE (OTC)

CHEMI SPA

* CHEMI SPA
 DECITABINE, DECITABINE
 TEMOZOLOMIDE, TEMOZOLOMIDE

CHEMISCH FBRK KRSSLR

* CHEMISCHE FABRIK KREUSSLER & CO. GMBH
 ASCLERA, POLIDOCANOL

CHEMO RESEARCH SL

* CHEMO RESEARCH SL
 BENZNIDAZOLE, BENZNIDAZOLE
 NUVESSA, METRONIDAZOLE

CHEPLAPHARM

* CHEPLAPHARM ARZNEIMITTEL GMBH
 CYTOVENE, GANCICLOVIR SODIUM
 ETOPOPHOS PRESERVATIVE FREE, ETOPOSIDE PHOSPHATE
 XENICAL, ORLISTAT

CHIA TAI TIANQING

* CHIA TAI TIANQING PHARMACEUTICAL GROUP CO LTD
 FULVESTRANT, FULVESTRANT

CHIESI USA INC

* 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
 CUROSURF, PORACTANT ALFA
 KENGREAL, CANGRELOR
 ZYFLO CR, ZILEUTON
 ZYFLO, ZILEUTON

CHILDRENS HOSP MI

* CHILDRENS HOSP MICHIGAN

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** C ****

* CHILDRENS HOSP MICHIGAN

FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18

CHINA RESOURCES

* CHINA RESOURCES SAIKE PHARMACEUTICAL CO LTD

AMLODIPINE BESYLATE, AMLODIPINE BESYLATE

CHIRHOCLIN

* CHIRHOCLIN INC

CHIRHOSTIM, SECRETIN SYNTHETIC HUMAN

CINTEX SVCS

* CINTEX SERVICES LLC

FLURANDRENOLIDE, FLURANDRENOLIDE

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

ALENDRONATE SODIUM, ALENDRONATE SODIUM

AMBRISENTAN, AMBRISENTAN

AMLODIPINE BESYLATE, AMLODIPINE BESYLATE

ANASTROZOLE, ANASTROZOLE

ATAZANAVIR SULFATE, ATAZANAVIR SULFATE

AZACITIDINE, AZACITIDINE

BIVALIRUDIN, BIVALIRUDIN

BLEOMYCIN SULFATE, BLEOMYCIN SULFATE

BOSENTAN, BOSENTAN

BUDESONIDE, BUDESONIDE

CELECOXIB, CELECOXIB

CINACALCET HYDROCHLORIDE, CINACALCET HYDROCHLORIDE

CYCLOPHOSPHAMIDE, CYCLOPHOSPHAMIDE

DARIFENACIN HYDROBROMIDE, DARIFENACIN HYDROBROMIDE

DECITABINE, DECITABINE

DEFERASIROX, DEFERASIROX

DICLOFENAC SODIUM, DICLOFENAC SODIUM

EFAVIRENZ, EFAVIRENZ

EFAVIRENZ, EMTRICITABINE, AND TENOFOVIR DISOPROXIL FUMARATE, EFAVIRENZ

EMTRICITABINE, EMTRICITABINE

ENTECAVIR, ENTECAVIR

EXEMESTANE, EXEMESTANE

FAMCICLOVIR, FAMCICLOVIR

FENOFIBRATE, FENOFIBRATE

FINASTERIDE, FINASTERIDE

FLUTAMIDE, FLUTAMIDE

GEMCITABINE HYDROCHLORIDE, GEMCITABINE HYDROCHLORIDE

GRISEOFULVIN, GRISEOFULVIN, MICROSIZE

ISOPROTERENOL HYDROCHLORIDE, ISOPROTERENOL HYDROCHLORIDE

LAMIVUDINE AND ZIDOVUDINE, LAMIVUDINE

LAMIVUDINE, LAMIVUDINE

LAMOTRIGINE, LAMOTRIGINE

LEVALBUTEROL HYDROCHLORIDE, LEVALBUTEROL HYDROCHLORIDE

MELOXICAM, MELOXICAM

METOPROLOL SUCCINATE, METOPROLOL SUCCINATE

MONTELUKAST SODIUM, MONTELUKAST SODIUM

NEVIRAPINE, NEVIRAPINE

OLOPATADINE HYDROCHLORIDE, OLOPATADINE HYDROCHLORIDE

OXALIPLATIN, OXALIPLATIN

PALONOSETRON HYDROCHLORIDE, PALONOSETRON HYDROCHLORIDE

PRAVASTATIN SODIUM, PRAVASTATIN SODIUM

PREGABALIN, PREGABALIN

RANOLAZINE, RANOLAZINE

SODIUM NITROPRUSSIDE, SODIUM NITROPRUSSIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

**** C ****

* CIPLA LTD
 SOLIFENACIN SUCCINATE, SOLIFENACIN SUCCINATE
 TADALAFIL, TADALAFIL
 TENOFOVIR DISOPROXIL FUMARATE, TENOFOVIR DISOPROXIL FUMARATE
 TERBINAFINE HYDROCHLORIDE, TERBINAFINE HYDROCHLORIDE
 TESTOSTERONE CYPIONATE, TESTOSTERONE CYPIONATE
 TESTOSTERONE, TESTOSTERONE
 TOPOTECAN HYDROCHLORIDE, TOPOTECAN HYDROCHLORIDE
 VALACYCLOVIR HYDROCHLORIDE, VALACYCLOVIR HYDROCHLORIDE
 VALGANCICLOVIR HYDROCHLORIDE, VALGANCICLOVIR HYDROCHLORIDE
 ZIDOVUDINE, ZIDOVUDINE
 ZOLEDRONIC ACID, ZOLEDRONIC ACID

CIPLA LTD

* CIPLA LTD
 ALBENDAZOLE, ALBENDAZOLE
 CARBOPLATIN, CARBOPLATIN
 CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
 EPIRUBICIN HYDROCHLORIDE, EPIRUBICIN HYDROCHLORIDE
 GRANISETRON HYDROCHLORIDE, GRANISETRON HYDROCHLORIDE
 IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE
 LEVOFLOXACIN, LEVOFLOXACIN
 TOPIRAMATE, TOPIRAMATE
 ZALEPLON, ZALEPLON
 ZIDOVUDINE, ZIDOVUDINE
 ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE

CIPLA USA

* CIPLA USA INC
 ZEMDRI, PLAZOMICIN SULFATE

CIRCASSIA

* CIRCASSIA LTD
 DUAKLIR PRESSAIR, ACLIDINIUM BROMIDE
 TUDORZA PRESSAIR, ACLIDINIUM BROMIDE

CLARUS

* CLARUS THERAPEUTICS INC
 JATENZO, TESTOSTERONE UNDECANOATE

CLINIGEN

* CLINIGEN INC
 ETHYOL, AMIFOSTINE
 TOTECT, DEXRAZOXANE HYDROCHLORIDE

CLINIGEN HLTHCARE

* CLINIGEN HEALTHCARE LTD
 FOSCAVIR, FOSCARNET SODIUM

CLIVUNEL INC

* CLINUVEL INC
 SCENESSE, AFAMELANOTIDE

CLOVER PHARMS

* CLOVER PHARMACEUTICALS CORP
 AMICAR, AMINOCAPROIC ACID

CLOVIS ONCOLOGY INC

* CLOVIS ONCOLOGY INC
 RUBRACA, RUCAPARIB CAMSYLATE

CMP DEV LLC

* CMP DEVELOPMENT LLC
 CAROSPIR, SPIRONOLACTONE
 POTASSIUM PHOSPHATES, POTASSIUM PHOSPHATE, DIBASIC

CMP PHARMA INC

* CMP PHARMA INC
 AMANTADINE HYDROCHLORIDE, AMANTADINE HYDROCHLORIDE
 HYDROCORTISONE IN ABSORBASE, HYDROCORTISONE
 ISONIAZID, ISONIAZID
 SODIUM POLYSTYRENE SULFONATE, SODIUM POLYSTYRENE SULFONATE
 SPS, SODIUM POLYSTYRENE SULFONATE
 TRIAMCINOLONE ACETONIDE IN ABSORBASE, TRIAMCINOLONE ACETONIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** C ******CNTY LINE PHARMS**

* COUNTY LINE PHARMACEUTICALS LLC
 BUTALBITAL AND ACETAMINOPHEN, ACETAMINOPHEN
 BUTALBITAL, ACETAMINOPHEN AND CAFFEINE, ACETAMINOPHEN
 CICLOPIROX, CICLOPIROX
 DYNACIN, MINOCYCLINE HYDROCHLORIDE
 FLUOCINONIDE, FLUOCINONIDE
 LIDEX, FLUOCINONIDE
 LIDEX-E, FLUOCINONIDE
 METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
 TAMBOCOR, FLECAINIDE ACETATE
 TRANDATE, LABETALOL HYDROCHLORIDE
 TRANLYCYPROMINE SULFATE, TRANLYCYPROMINE SULFATE
 UREX, METHENAMINE HIPPURATE

CODY LABS INC

* CODY LABORATORIES INC A WHOLLY OWNED SUBSIDIARY OF LANNETT CO INC
 NUMBRINO, COCAINE HYDROCHLORIDE

COEPTIS

* COEPTIS PHARMACEUTICALS INC
 CONSENSI, AMLODIPINE BESYLATE

COLGATE PALMOLIVE CO

* COLGATE PALMOLIVE CO
 PERIOGARD, CHLORHEXIDINE GLUCONATE

COLGATE-PALMOLIVE CO

* COLGATE-PALMOLIVE CO
 PERIOGARD, CHLORHEXIDINE GLUCONATE

COLLEGIUM PHARM INC

* COLLEGIUM PHARMACEUTICAL INC
 NUCYNTA ER, TAPENTADOL HYDROCHLORIDE
 NUCYNTA, TAPENTADOL HYDROCHLORIDE
 XTAMPZA ER, OXYCODONE

COMBE

* COMBE INC
 VAGISTAT-1, TIOCONAZOLE (OTC)

CONCORD BIOTECH LTD

* CONCORD BIOTECH LTD
 MYCOPHENOLATE MOFETIL, MYCOPHENOLATE MOFETIL
 MYCOPHENOLIC ACID, MYCOPHENOLIC ACID

CONCORDIA

* CONCORDIA PHARMACEUTICALS INC
 DUTOPROL, HYDROCHLOROTHIAZIDE
 DYRENIUM, TRIAMTERENE
 LANOXIN, DIGOXIN
 NILANDRON, NILUTAMIDE
 PANRETIN, ALITRETINOIN
 PARNATE, TRANLYCYPROMINE SULFATE
 PLAQUENIL, HYDROXYCHLOROQUINE SULFATE
 SALAGEN, PILOCARPINE HYDROCHLORIDE
 UROXATRAL, ALFUZOSIN HYDROCHLORIDE

CONCORDIA PHARMS INC

* CONCORDIA PHARMACEUTICALS INC
 KAPVAY, CLONIDINE HYDROCHLORIDE
 ORAPRED ODT, PREDNISOLONE SODIUM PHOSPHATE

CONTRACT PHARMACAL

* CONTRACT PHARMACAL CORP
 CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 IBUPROFEN, IBUPROFEN
 IBUPROFEN, IBUPROFEN (OTC)
 NAPROXEN SODIUM, NAPROXEN SODIUM (OTC)

COOPERSURGICAL

* COOPERSURGICAL INC
 PARAGARD T 380A, COPPER

CORCEPT THERAP

* CORCEPT THERAPEUTICS INC

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** C ****

* CORCEPT THERAPEUTICS INC
KORLYM, MIFEPRISTONE

CORDEN PHARMA

* CORDEN PHARMA LATINA SPA
GLEOSTINE, LOMUSTINE

COREPHARMA

* COREPHARMA LLC
BENAZEPRIL HYDROCHLORIDE, BENAZEPRIL HYDROCHLORIDE
EPLERENONE, EPLERENONE
LISINOPRIL, LISINOPRIL
LOVASTATIN, LOVASTATIN
SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
TORSEMIDE, TORSEMIDE

COSETTE

* COSETTE PHARMACEUTICALS INC
ALBUTEROL SULFATE, ALBUTEROL SULFATE
DESOXIMETASONE, DESOXIMETASONE
HYDROCORTISONE VALERATE, HYDROCORTISONE VALERATE
MIGERGOT, CAFFEINE

COVIS PHARMA BV

* COVIS PHARMA BV
ALTOPREV, LOVASTATIN
ALVESCO, CICLESONIDE
BETAPACE AF, SOTALOL HYDROCHLORIDE
BETAPACE, SOTALOL HYDROCHLORIDE
LANOXIN PEDIATRIC, DIGOXIN
LANOXIN, DIGOXIN
OMNARIS, CICLESONIDE
PRILOSEC, OMEPRAZOLE MAGNESIUM
RILUTEK, RILUZOLE
SULAR, NISOLDIPINE
ZANAFLEX, TIZANIDINE HYDROCHLORIDE
ZETONNA, CICLESONIDE

CPDC

* CENTRE FOR PROBE DEVELOPMENT AND COMMERCIALIZATION
FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18

CPPI CV

* CP PHARMACEUTICALS INTERNATIONAL CV
SUTENT, SUNITINIB MALATE

CROSSMEDIKA SA

* CROSSMEDIKA SA
MOXIFLOXACIN HYDROCHLORIDE, MOXIFLOXACIN HYDROCHLORIDE
TRIMIPRAMINE MALEATE, TRIMIPRAMINE MALEATE
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

CSPC OUYI

* CSPC OUYI PHARMACEUTICAL CO LTD
AZITHROMYCIN, AZITHROMYCIN
BENZONATATE, BENZONATATE
CELECOXIB, CELECOXIB
CLOPIDOGREL BISULFATE, CLOPIDOGREL BISULFATE
CONJUPRI, LEVAMLODIPINE MALEATE
DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE
GABAPENTIN, GABAPENTIN
MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** C ****

* CSPC OUYI PHARMACEUTICAL CO LTD
 MONTELUKAST SODIUM, MONTELUKAST SODIUM
 PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE
 PREGABALIN, PREGABALIN
 SOLIFENACIN SUCCINATE, SOLIFENACIN SUCCINATE

CSPC OUYI PHARM CO

* CSPC OUYI PHARMACEUTICAL CO LTD
 TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE

CUBIST PHARMS

* CUBIST PHARMACEUTICALS INC
 ENTEREG, ALVIMOPAN

CUBIST PHARMS LLC

* CUBIST PHARMACEUTICALS LLC
 CUBICIN RF, DAPTOMYCIN
 CUBICIN, DAPTOMYCIN
 DIFICID, FIDAXOMICIN
 SIVEXTRO, TEDIZOLID PHOSPHATE
 ZERBAXA, CEFTOLOZANE SULFATE

CUMBERLAND PHARMS

* CUMBERLAND PHARMACEUTICALS INC
 ACETADOTE, ACETYLCYSTEINE
 CALDOLOR, IBUPROFEN
 LACTULOSE, LACTULOSE
 OMEPRAZOLE AND CLARITHROMYCIN AND AMOXICILLIN, AMOXICILLIN
 REDITREX, METHOTREXATE
 VAPRISOL IN 5% DEXTROSE IN PLASTIC CONTAINER, CONIVAPTAN HYDROCHLORIDE
 VIBATIV, TELAVANCIN HYDROCHLORIDE

CURIUM

* CURIUM US LLC
 GALLIUM CITRATE GA 67, GALLIUM CITRATE GA-67
 INDIUM IN 111 CHLORIDE, INDIUM IN-111 CHLORIDE
 OCTREOSCAN, INDIUM IN-111 PENTETREOTIDE KIT
 SODIUM IODIDE I 123, SODIUM IODIDE I-123
 TECHNESCAN MAG3, TECHNETIUM TC-99M MERTIATIDE KIT
 TECHNESCAN PYP KIT, TECHNETIUM TC-99M PYROPHOSPHATE KIT
 TECHNESCAN, 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
 ACETAMINOPHEN, ACETAMINOPHEN
 DICYCLOMINE HYDROCHLORIDE, DICYCLOMINE HYDROCHLORIDE
 DIHYDROERGOTAMINE MESYLATE, DIHYDROERGOTAMINE MESYLATE
 FLUDARABINE PHOSPHATE, FLUDARABINE PHOSPHATE
 PENTOBARBITAL SODIUM, PENTOBARBITAL SODIUM
 SODIUM TETRADECYL SULFATE, SODIUM TETRADECYL SULFATE
 VALRUBICIN, VALRUBICIN

CYCLE PHARMS LTD

* CYCLE PHARMACEUTICALS LTD
 KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
 NITYR, NITISINONE

**** D ******DAEWOONG PHARM CO**

* DAEWOONG PHARMACEUTICAL CO LTD
 MEROPENEM, MEROPENEM

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** D ******DAIICHI SANKYO**

* DAIICHI SANKYO INC
 AZOR, AMLODIPINE BESYLATE
 BENICAR HCT, HYDROCHLOROTHIAZIDE
 BENICAR, OLMESARTAN MEDOXOMIL
 TRIBENZOR, AMLODIPINE BESYLATE
 WELCHOL, COLESEVELAM HYDROCHLORIDE

DAIICHI SANKYO INC

* DAIICHI SANKYO INC
 EVOXAC, CEVIMELINE HYDROCHLORIDE
 MORPHABOND ER, MORPHINE SULFATE
 SAVAYSA, EDOXABAN TOSYLATE
 TURALIO, PEXIDARTINIB HYDROCHLORIDE
 WELCHOL, COLESEVELAM HYDROCHLORIDE

DANCO LABS LLC

* DANCO LABORATORIES LLC
 MIFEPREX, MIFEPRISTONE

DAVA PHARMS INC

* DAVA PHARMACEUTICALS INC
 ACYCLOVIR, ACYCLOVIR
 AMOXICILLIN, AMOXICILLIN
 AMPICILLIN TRIHYDRATE, AMPICILLIN/AMPICILLIN TRIHYDRATE
 ATENOLOL, ATENOLOL
 CIMETIDINE HYDROCHLORIDE, CIMETIDINE HYDROCHLORIDE
 DOXAZOSIN MESYLATE, DOXAZOSIN MESYLATE
 GLYBURIDE (MICRONIZED), GLYBURIDE
 METHOTREXATE SODIUM, METHOTREXATE SODIUM
 MORPHINE SULFATE, MORPHINE SULFATE
 PENICILLIN V POTASSIUM, PENICILLIN V POTASSIUM
 PROPYLTHIOURACIL, PROPYLTHIOURACIL
 VOSPIRE ER, ALBUTEROL SULFATE

DAVIS AND GECK

* DAVIS AND GECK DIV AMERICAN CYANAMID CO
 PRE-OP II, HEXACHLOROPHENE
 PRE-OP, HEXACHLOROPHENE

DBL PHARMS

* DBL PHARMACEUTICALS INC
 METHOCARBAMOL, METHOCARBAMOL

DECATUR

* DECATUR MEMORIAL HOSP
 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

DENTSPLY PHARM

* DENTSPLY PHARMACEUTICAL INC
 CITANEST FORTE DENTAL, EPINEPHRINE BITARTRATE
 ORAQIX, LIDOCAINE

DEPROCO

* DEPROCO INC
 LIGNOSPAN FORTE, EPINEPHRINE BITARTRATE
 LIGNOSPAN STANDARD, EPINEPHRINE BITARTRATE
 SCANDONEST L, LEVONORDEFRIN
 SCANDONEST PLAIN, MEPIVACAINE HYDROCHLORIDE
 SEPTOCAINE, ARTICAINE HYDROCHLORIDE

DERMIRA INC

* DERMIRA INC
 QBREXZA, GLYCOPYRRONIUM TOSYLATE

DEVA HOLDING AS

* DEVA HOLDING AS
 ESOMEPRAZOLE SODIUM, ESOMEPRAZOLE SODIUM
 TEMOZOLOMIDE, TEMOZOLOMIDE

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
 DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
 HEMADY, DEXAMETHASONE
 LANSOPRAZOLE, LANSOPRAZOLE (OTC)
 LEVETIRACETAM, LEVETIRACETAM
 OMEPRAZOLE, OMEPRAZOLE (OTC)
 PERIOCHIP, CHLORHEXIDINE GLUCONATE
 VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE

DFB ONCOLOGY LTD

* DFB ONCOLOGY LTD
 DOCETAXEL, DOCETAXEL

DIAGNOSTIC GREEN

* DIAGNOSTIC GREEN GMBH
 INDOCYANINE GREEN, INDOCYANINE GREEN

DIALYSIS SUPS

* DIALYSIS SUPPLIES INC
 NORMOCARB HF 25, MAGNESIUM CHLORIDE
 NORMOCARB HF 35, MAGNESIUM CHLORIDE

DIGESTIVE CARE INC

* DIGESTIVE CARE INC
 PERTZYE, PANCRELIPASE (AMYLASE)

DORC

* DORC INTERNATIONAL BV
 MEMBRANEBLUE, TRYPAN BLUE
 VISIONBLUE, TRYPAN BLUE

DOUGLAS PHARMS

* DOUGLAS PHARMACEUTICALS AMERICA LTD
 MYORISAN, ISOTRETINOIN

DOW PHARM

* DOW PHARMACEUTICAL SCIENCES
 ALTRENO, TRETINOIN
 ATRALIN, TRETINOIN

DR REDDYS

* DR REDDYS LABORATORIES INC
 NITROGLYCERIN, NITROGLYCERIN
 PROGESTERONE, PROGESTERONE
 PROPOFOL, PROPOFOL
 SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
 TESTOSTERONE, TESTOSTERONE
 ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE

* DR REDDYS LABORATORIES LTD
 AMLODIPINE BESYLATE AND ATORVASTATIN CALCIUM, AMLODIPINE BESYLATE
 ASPIRIN AND DIPYRIDAMOLE, ASPIRIN
 AZACITIDINE, AZACITIDINE
 CARFILZOMIB, CARFILZOMIB
 COLCHICINE, COLCHICINE
 COLESEVELAM HYDROCHLORIDE, COLESEVELAM HYDROCHLORIDE
 DAPTOMYCIN, DAPTOMYCIN
 DECITABINE, DECITABINE
 DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
 ENALAPRILAT, ENALAPRILAT
 GRANISETRON HYDROCHLORIDE, GRANISETRON HYDROCHLORIDE
 IMATINIB MESYLATE, IMATINIB MESYLATE
 NALOXONE HYDROCHLORIDE, NALOXONE HYDROCHLORIDE
 NEOSTIGMINE METHYLSULFATE, NEOSTIGMINE METHYLSULFATE
 NYSTATIN AND TRIAMCINOLONE ACETONIDE, NYSTATIN
 OMEPRAZOLE, OMEPRAZOLE (OTC)
 PAMIDRONATE DISODIUM, PAMIDRONATE DISODIUM
 PANCURONIUM BROMIDE, PANCURONIUM BROMIDE
 PARICALCITOL, PARICALCITOL
 PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

**** D ****

- * DR REDDYS LABORATORIES LTD
 - PREGABALIN, PREGABALIN
 - RABEPRAZOLE SODIUM, RABEPRAZOLE SODIUM
 - SUCCINYLCHOLINE CHLORIDE, SUCCINYLCHOLINE CHLORIDE
 - TETRABENAZINE, TETRABENAZINE
 - THIOTEPA, THIOTEPA
 - VALGANCICLOVIR HYDROCHLORIDE, VALGANCICLOVIR HYDROCHLORIDE
 - VINORELBINE TARTRATE, VINORELBINE TARTRATE

DR REDDYS LA

- * DR REDDYS LABORATORIES LOUISIANA LLC
 - IBUPROFEN, IBUPROFEN
 - IBUPROFEN, IBUPROFEN (OTC)
 - LOPURIN, ALLOPURINOL
 - SSD, SILVER SULFADIAZINE

DR REDDYS LABS INC

- * DR REDDYS LABORATORIES INC
 - AMLODIPINE BESYLATE AND BENAZEPRIL HYDROCHLORIDE, AMLODIPINE BESYLATE
 - CLOPIDOGREL BISULFATE, CLOPIDOGREL BISULFATE
 - 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
 - ATOMOXETINE HYDROCHLORIDE, ATOMOXETINE HYDROCHLORIDE
 - ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM
 - BIVALIRUDIN, BIVALIRUDIN
 - BORTEZOMIB, BORTEZOMIB
 - CANDESARTAN CILEXETIL AND HYDROCHLOROTHIAZIDE, CANDESARTAN CILEXETIL
 - CARBOPROST TROMETHAMINE, CARBOPROST TROMETHAMINE
 - 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
 - CLOFARABINE, CLOFARABINE
 - CLOPIDOGREL BISULFATE, CLOPIDOGREL BISULFATE
 - DESLORATADINE AND PSEUDOEPHEDRINE SULFATE 24 HOUR, DESLORATADINE
 - DESLORATADINE, DESLORATADINE
 - DIVALPROEX SODIUM, DIVALPROEX SODIUM
 - DOCETAXEL, DOCETAXEL
 - DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
 - DOXORUBICIN HYDROCHLORIDE (LIPOSOMAL), DOXORUBICIN HYDROCHLORIDE
 - ENALAPRIL MALEATE AND HYDROCHLOROTHIAZIDE, ENALAPRIL MALEATE
 - ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM
 - ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM (OTC)
 - ESZOPICLONE, ESZOPICLONE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** D ****

* DR REDDYS LABORATORIES LTD
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
GLYCOPYRROLATE, GLYCOPYRROLATE
GRANISETRON HYDROCHLORIDE, GRANISETRON HYDROCHLORIDE
GUAIFENESIN AND PSEUDOEPHEDRINE HYDROCHLORIDE, GUAIFENESIN (OTC)
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)
LATANOPROST, LATANOPROST
LETROZOLE, LETROZOLE
LEVETIRACETAM, LEVETIRACETAM
LEVOCETIRIZINE DIHYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE (OTC)
MELPHALAN HYDROCHLORIDE, MELPHALAN HYDROCHLORIDE
MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
METOPROLOL SUCCINATE, METOPROLOL SUCCINATE
MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE
MONTELUKAST SODIUM, MONTELUKAST SODIUM
MOXIFLOXACIN HYDROCHLORIDE, MOXIFLOXACIN HYDROCHLORIDE
NAPROXEN AND ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM
NAPROXEN SODIUM, NAPROXEN SODIUM
NATEGLINIDE, NATEGLINIDE
NIZATIDINE, NIZATIDINE
OFLOXACIN, OFLOXACIN
OLANZAPINE, OLANZAPINE
OMEPRAZOLE AND SODIUM BICARBONATE, OMEPRAZOLE
OMEPRAZOLE MAGNESIUM, OMEPRAZOLE MAGNESIUM (OTC)
OMEPRAZOLE, OMEPRAZOLE
ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
OXAPROZIN, OXAPROZIN
PALONOSETRON HYDROCHLORIDE, PALONOSETRON HYDROCHLORIDE
PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
PARICALCITOL, PARICALCITOL
PHYTONADIONE, PHYTONADIONE
QUETIAPINE FUMARATE, QUETIAPINE FUMARATE
RAMIPRIL, RAMIPRIL
RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE (OTC)
RISPERIDONE, RISPERIDONE
ROPINIROLE HYDROCHLORIDE, ROPINIROLE HYDROCHLORIDE
SEVELAMER CARBONATE, SEVELAMER CARBONATE
SIROLIMUS, SIROLIMUS
SODIUM NITROPRUSSIDE, SODIUM NITROPRUSSIDE
TACROLIMUS, TACROLIMUS
TADALAFIL, TADALAFIL
TOPOTECAN HYDROCHLORIDE, TOPOTECAN HYDROCHLORIDE
TRIENTINE HYDROCHLORIDE, TRIENTINE HYDROCHLORIDE
VALGANCICLOVIR HYDROCHLORIDE, VALGANCICLOVIR HYDROCHLORIDE
VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE
VIGABATRIN, VIGABATRIN
ZAFIRLUKAST, ZAFIRLUKAST

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** D ****

* DR REDDYS LABORATORIES LTD
ZENATANE, ISOTRETINOIN
ZOLEDRONIC ACID, ZOLEDRONIC ACID

DR REDDYS LABS SA

* DR REDDYS LABORATORIES SA
BUPRENORPHINE HYDROCHLORIDE AND NALOXONE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE
EZETIMIBE AND SIMVASTATIN, EZETIMIBE
FENOFIBRATE (MICRONIZED), FENOFIBRATE
HABITROL, NICOTINE (OTC)
MERCAPTOPYRINE, MERCAPTOPYRINE
NICOTINE POLACRILEX, NICOTINE POLACRILEX (OTC)
RAMELTEON, RAMELTEON
TOBRAMYCIN, TOBRAMYCIN

DRAXIMAGE

* DRAXIMAGE INC
TECHNETIUM TC 99M ALBUMIN AGGREGATED KIT, TECHNETIUM TC-99M ALBUMIN AGGREGATED KIT

DUCHESNAY

* DUCHESNAY INC
BONJESTA, DOXYLAMINE SUCCINATE
DICLEGIS, DOXYLAMINE SUCCINATE
OSPHENA, OSPEMIFENE

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
METHYLPREDNISOLONE, METHYLPREDNISOLONE
TRIAMTERENE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
VELIVET, DESOGESTREL

DUSA

* DUSA PHARMACEUTICALS INC
LEVULAN, AMINOLEVULINIC ACID HYDROCHLORIDE

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

* EAGLE PHARMACEUTICALS INC
ARGATROBAN IN SODIUM CHLORIDE, ARGATROBAN
BELRAPZO, BENDAMUSTINE HYDROCHLORIDE
BENDEKA, BENDAMUSTINE HYDROCHLORIDE
PEMFEXY, PEMETREXED
RYANODEX, DANTROLENE SODIUM

ECI PHARMS LLC

* ECI PHARMACEUTICALS LLC
BETHANECHOL CHLORIDE, BETHANECHOL CHLORIDE
HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE
LAMIVUDINE, LAMIVUDINE
LEVETIRACETAM, LEVETIRACETAM
METHIMAZOLE, METHIMAZOLE
RIZATRIPTAN BENZOATE, RIZATRIPTAN BENZOATE
SODIUM POLYSTYRENE SULFONATE, SODIUM POLYSTYRENE SULFONATE
VALPROIC ACID, VALPROIC ACID

ECOLAB

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** E ****

* ECOLAB INC
 CHG SCRUB, CHLORHEXIDINE GLUCONATE (OTC)
 CIDA-STAT, CHLORHEXIDINE GLUCONATE (OTC)

ECR

* ECR PHARMACEUTICALS
 DEXAMETHASONE, DEXAMETHASONE

ECR PHARMA

* ECR PHARMA
 TUSSICAPS, CHLORPHENIRAMINE POLISTIREX

EDENBRIDGE PHARMS

* EDENBRIDGE PHARMACEUTICALS LLC
 ALBENDAZOLE, ALBENDAZOLE
 CARBIDOPA, CARBIDOPA
 DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
 ETHACRYNIC ACID, ETHACRYNIC ACID
 ETODOLAC, ETODOLAC
 IVERMECTIN, IVERMECTIN
 PREDNISOLONE SODIUM PHOSPHATE, PREDNISOLONE SODIUM PHOSPHATE
 TINIDAZOLE, TINIDAZOLE

EDISON THERAPS LLC

* EDISON THERAPEUTICS LLC
 METHERGINE, METHYLERGONOVINE MALEATE

EGALET

* EGALET US INC
 INDOCIN, INDOMETHACIN

EI INC

* EI INC
 THEROXIDIL, MINOXIDIL (OTC)

EISAI INC

* EISAI INC
 ACIPHEX, RABEPRAZOLE SODIUM
 ARICEPT, DONEPEZIL HYDROCHLORIDE
 BANZEL, RUFINAMIDE
 FYCOMPA, PERAMPANEL
 HALAVEN, ERIBULIN MESYLATE
 LENVIMA, LENVATINIB MESYLATE

ELI LILLY AND CO

* ELI LILLY AND CO
 BAQSIMI, GLUCAGON
 BASAGLAR, INSULIN GLARGINE
 EFFIENT, PRASUGREL HYDROCHLORIDE
 HUMALOG KWIKPEN, INSULIN LISPRO RECOMBINANT
 OLUMIANT, BARICITINIB
 PROZAC, FLUOXETINE HYDROCHLORIDE
 REYVOW, LASMIDITAN SUCCINATE
 VERZENIO, ABEMACICLIB

ELI LILLY CO

* ELI LILLY CO
 ADCIRCA, TADALAFIL
 ZYPREXA RELPREVV, OLANZAPINE PAMOATE

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
 ISRADIPINE, ISRADIPINE
 LOXAPINE SUCCINATE, LOXAPINE SUCCINATE
 METHADONE HYDROCHLORIDE, METHADONE HYDROCHLORIDE
 PHENDIMETRAZINE TARTRATE, PHENDIMETRAZINE TARTRATE
 PHENTERMINE HYDROCHLORIDE, PHENTERMINE HYDROCHLORIDE
 TRIMIPRAMINE MALEATE, TRIMIPRAMINE MALEATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** E ******ELYSIUM**

* ELYSIUM PHARMACEUTICALS LTD
 CALCITRIOL, CALCITRIOL
 CYPROHEPTADINE HYDROCHLORIDE, CYPROHEPTADINE HYDROCHLORIDE
 PYRIDOSTIGMINE BROMIDE, PYRIDOSTIGMINE BROMIDE
 TIMOLOL MALEATE, TIMOLOL MALEATE

EMCURE PHARMS

* EMCURE PHARMACEUTICALS LTD
 ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE

EMCURE PHARMS LTD

* EMCURE PHARMACEUTICALS LTD
 ACARBOSE, ACARBOSE
 AMIKACIN SULFATE, AMIKACIN SULFATE
 BENZPHETAMINE HYDROCHLORIDE, BENZPHETAMINE HYDROCHLORIDE
 BICNU, CARMUSTINE
 CIDOFOVIR, CIDOFOVIR
 COLISTIMETHATE SODIUM, COLISTIMETHATE SODIUM
 DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
 ETOMIDATE, ETOMIDATE
 FOSINOPRIL SODIUM AND HYDROCHLOROTHIAZIDE, FOSINOPRIL SODIUM
 FUROSEMIDE, FUROSEMIDE
 GEMCITABINE HYDROCHLORIDE, GEMCITABINE HYDROCHLORIDE
 IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE
 METOCLOPRAMIDE, METOCLOPRAMIDE HYDROCHLORIDE
 ONDANSETRON HYDROCHLORIDE PRESERVATIVE FREE, ONDANSETRON HYDROCHLORIDE
 OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE
 PROCHLORPERAZINE EDISYLATE, PROCHLORPERAZINE EDISYLATE
 RIFAMPIN, RIFAMPIN
 RIZATRIPTAN BENZOATE, RIZATRIPTAN BENZOATE
 TRANEXAMIC ACID, TRANEXAMIC ACID
 ZOLEDRONIC ACID, ZOLEDRONIC ACID

EMD SERONO

* EMD SERONO INC
 GONAL-F RFF REDI-JECT, FOLLITROPIN ALFA/BETA
 GONAL-F RFF, FOLLITROPIN ALFA/BETA
 GONAL-F, FOLLITROPIN ALFA/BETA
 OVIDREL, CHORIOGONADOTROPIN ALFA
 SAIZEN, SOMATROPIN
 SEROSTIM, SOMATROPIN
 ZORBTIVE, SOMATROPIN RECOMBINANT

EMD SERONO INC

* EMD SERONO INC
 CETROTIDE, CETRORELIX
 MAVENCLAD, CLADRIBINE

EMERALD INTL LTD

* EMERALD INTERNATIONAL LTD
 BACLOFEN, BACLOFEN

EMMAUS MEDCL

* EMMAUS MEDICAL INC
 ENDARI, L-GLUTAMINE

ENCORE DERMAT

* ENCORE DERMATOLOGY INC
 IMPOYZ, CLOBETASOL PROPIONATE
 SERNIVO, BETAMETHASONE DIPROPIONATE

ENCUBE

* ENCUBE ETHICALS PRIVATE LTD
 CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
 DICLOFENAC SODIUM, DICLOFENAC SODIUM
 TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE

ENCUBE ETHICALS

* ENCUBE ETHICALS PVT LTD
 CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
 DESONIDE, DESONIDE
 FLUOCINOLONE ACETONIDE, FLUOCINOLONE ACETONIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

**** E ****

* ENCUBE ETHICALS PVT LTD
 FLUOCINONIDE, FLUOCINONIDE
 PERMETHRIN, PERMETHRIN

ENDO PHARM

* ENDO PHARMACEUTICAL SOLUTIONS INC
 SUPPRELIN LA, HISTRELIN ACETATE
 VALSTAR PRESERVATIVE FREE, VALRUBICIN
 VANTAS, HISTRELIN ACETATE

ENDO PHARMS

* ENDO PHARMACEUTICALS INC
 FORTESTA, TESTOSTERONE
 FROVA, FROVATRIPTAN SUCCINATE
 PERCODAN, ASPIRIN

ENDO PHARMS INC

* ENDO PHARMACEUTICALS INC
 AVEED, TESTOSTERONE UNDECANOATE
 COLY-MYCIN S, COLISTIN SULFATE
 MEGACE ES, MEGESTROL ACETATE
 NASCOBAL, CYANOCOBALAMIN
 VALGANCICLOVIR HYDROCHLORIDE, VALGANCICLOVIR HYDROCHLORIDE

EPI HLTH

* EPI HEALTH LLC
 CLODERM, CLOCORTOLONE PIVALATE
 MINOLIRA, MINOCYCLINE HYDROCHLORIDE
 RHOFAD, OXYMETAZOLINE HYDROCHLORIDE
 SITAVIG, ACYCLOVIR

EPIC PHARMA

* EPIC PHARMA INC
 MEPERIDINE HYDROCHLORIDE, MEPERIDINE HYDROCHLORIDE
 NICARDIPINE HYDROCHLORIDE, NICARDIPINE HYDROCHLORIDE

* EPIC PHARMA LLC
 BETAXOLOL HYDROCHLORIDE, BETAXOLOL HYDROCHLORIDE
 CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE
 FLAVOXATE HYDROCHLORIDE, FLAVOXATE HYDROCHLORIDE
 SULINDAC, SULINDAC
 TRANDOLAPRIL, TRANDOLAPRIL
 URSODIOL, URSODIOL

EPIC PHARMA INC

* EPIC PHARMA INC
 ESTRADIOL, ESTRADIOL
 SOTALOL HYDROCHLORIDE, SOTALOL HYDROCHLORIDE

EPIC PHARMA LLC

* EPIC PHARMA LLC
 AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
 AZITHROMYCIN, AZITHROMYCIN
 BENZPHETAMINE HYDROCHLORIDE, BENZPHETAMINE HYDROCHLORIDE
 BENZTROPINE MESYLATE, BENZTROPINE MESYLATE
 BUSPIRONE HYDROCHLORIDE, BUSPIRONE HYDROCHLORIDE
 CLINDAMYCIN HYDROCHLORIDE, CLINDAMYCIN HYDROCHLORIDE
 DEMECLOCYCLINE HYDROCHLORIDE, DEMECLOCYCLINE HYDROCHLORIDE
 DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE
 DOXYCYCLINE HCLATE, DOXYCYCLINE HCLATE
 GABAPENTIN, GABAPENTIN
 GLIPIZIDE AND METFORMIN HYDROCHLORIDE, GLIPIZIDE
 GLYBURIDE, GLYBURIDE
 GUANFACINE HYDROCHLORIDE, GUANFACINE HYDROCHLORIDE
 HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
 HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE
 MECLIZINE HYDROCHLORIDE, MECLIZINE HYDROCHLORIDE
 METHADONE HYDROCHLORIDE, METHADONE HYDROCHLORIDE
 MOLINDONE HYDROCHLORIDE, MOLINDONE HYDROCHLORIDE
 NYSTATIN, NYSTATIN
 OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN
 OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
 OXYMORPHONE HYDROCHLORIDE, OXYMORPHONE HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

**** E ****

* EPIC PHARMA LLC
 PHENYTOIN, PHENYTOIN
 POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
 PROTRIPTYLINE HYDROCHLORIDE, PROTRIPTYLINE HYDROCHLORIDE
 SODIUM POLYSTYRENE SULFONATE, SODIUM POLYSTYRENE SULFONATE
 TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE

EPIZYME INC

* EPIZYME INC
 TAZVERIK, TAZEMETOSTAT HYDROBROMIDE

ESPERION THERAPS INC

* ESPERION THERAPEUTICS INC
 NEXLETOL, BEMPEDOIC ACID
 NEXLIZET, BEMPEDOIC ACID

ESPERO

* ESPERO BIOPHARMA INC
 DURLAZA, ASPIRIN

ESSENTIAL ISOTOPES

* ESSENTIAL ISOTOPES LLC
 AMMONIA N 13, AMMONIA N-13
 FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18
 SODIUM FLUORIDE F-18, SODIUM FLUORIDE F-18

ETHYPHARM

* ETHYPHARM
 BUPRENORPHINE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE

ETHYPHARM USA CORP

* ETHYPHARM USA CORP
 BUPRENORPHINE HYDROCHLORIDE AND NALOXONE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE

ETON PHARMS

* ETON PHARMACEUTICALS
 BIORPHEN, PHENYLEPHRINE HYDROCHLORIDE

EUGIA PHARMA

* EUGIA PHARMA SPECIALITIES LTD
 CAPECITABINE, CAPECITABINE
 CARBOPLATIN, CARBOPLATIN
 HYDROXYPROGESTERONE CAPROATE, HYDROXYPROGESTERONE CAPROATE
 LETROZOLE, LETROZOLE
 METHOTREXATE SODIUM, METHOTREXATE SODIUM
 OXALIPLATIN, OXALIPLATIN
 PROGESTERONE, PROGESTERONE

EUROHLTH INTL SARL

* EUROHEALTH INTERNATIONAL SARL
 DROPERIDOL, DROPERIDOL
 HYDROMORPHONE HYDROCHLORIDE, HYDROMORPHONE HYDROCHLORIDE
 NEOSTIGMINE METHYLSULFATE, NEOSTIGMINE METHYLSULFATE

EVUS

* EVUS HEALTH SOLUTIONS LLC
 NITROMIST, NITROGLYCERIN

EXALENZ BIOSCIENCE

* EXALENZ BIOSCIENCE LTD
 IDKIT:HP, CITRIC ACID

EXELA PHARMA SCIENCE

* EXELA PHARMA SCIENCES
 CAFFEINE CITRATE, CAFFEINE CITRATE
 NICARDIPINE HYDROCHLORIDE, NICARDIPINE HYDROCHLORIDE

EXELA PHARMA SCS LLC

* EXELA PHARMA SCIENCES LLC
 CAFFEINE CITRATE, CAFFEINE CITRATE
 ELCYS, CYSTEINE HYDROCHLORIDE
 GANZYK-RTU, GANCICLOVIR
 GLYRX-PF, GLYCOPYRROLATE
 MAGNESIUM SULFATE, MAGNESIUM SULFATE
 NIPRIDE RTU IN SODIUM CHLORIDE 0.9%, SODIUM NITROPRUSSIDE
 POTASSIUM ACETATE, POTASSIUM ACETATE
 SODIUM BICARBONATE, SODIUM BICARBONATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

**** E ****

* EXELA PHARMA SCIENCES LLC
 TRANEXAMIC ACID, TRANEXAMIC ACID
 VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE

EXELIXIS

* EXELIXIS INC
 COMETRIQ, CABOZANTINIB S-MALATE

EXELIXIS INC

* EXELIXIS INC
 CABOMETYX, CABOZANTINIB S-MALATE

EXELTIS USA INC

* EXELTIS USA INC
 SLYND, DROSPIRENONE

EYEPOINT PHARMS

* EYEPOINT PHARMACEUTICALS INC
 DEXYCU KIT, DEXAMETHASONE
 YUTIQ, FLUOCINOLONE ACETONIDE

EYEVANCE

* EYEVANCE PHARMACEUTICALS LLC
 FLAREX, FLUOROMETHOLONE ACETATE
 TOBRADEX ST, DEXAMETHASONE
 ZERVIAE, CETIRIZINE HYDROCHLORIDE

EYWA

* EYWA PHARMA INC
 ACETAMINOPHEN AND CODEINE PHOSPHATE, ACETAMINOPHEN
 OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE

EYWA PHARMA

* EYWA PHARMA PTE LTD
 ACETAZOLAMIDE, ACETAZOLAMIDE
 ALOSETRON HYDROCHLORIDE, ALOSETRON HYDROCHLORIDE
 BACLOFEN, BACLOFEN
 URSODIOL, URSODIOL

LILLY

* ELI LILLY AND CO
 ALIMTA, PEMETREXED DISODIUM
 CIALIS, TADALAFIL
 CYMBALTA, DULOXETINE HYDROCHLORIDE
 EVISTA, RALOXIFENE HYDROCHLORIDE
 FORTEO, TERIPARATIDE
 GEMZAR, GEMCITABINE HYDROCHLORIDE
 GLUCAGON, GLUCAGON
 HUMALOG KWIKPEN, INSULIN LISPRO RECOMBINANT
 HUMALOG MIX 50/50 KWIKPEN, INSULIN LISPRO PROTAMINE RECOMBINANT
 HUMALOG MIX 50/50, INSULIN LISPRO PROTAMINE RECOMBINANT
 HUMALOG MIX 75/25 KWIKPEN, INSULIN LISPRO PROTAMINE RECOMBINANT
 HUMALOG MIX 75/25, INSULIN LISPRO PROTAMINE RECOMBINANT
 HUMALOG TEMPO PEN, INSULIN LISPRO RECOMBINANT
 HUMALOG, INSULIN LISPRO RECOMBINANT
 HUMATROPE, SOMATROPIN
 HUMULIN 70/30 PEN, INSULIN RECOMBINANT HUMAN (OTC)
 HUMULIN 70/30, INSULIN RECOMBINANT HUMAN (OTC)
 HUMULIN N, INSULIN SUSP ISOPHANE RECOMBINANT HUMAN (OTC)
 HUMULIN R KWIKPEN, INSULIN HUMAN
 HUMULIN R PEN, INSULIN RECOMBINANT HUMAN (OTC)
 HUMULIN R, INSULIN HUMAN
 HUMULIN R, INSULIN RECOMBINANT HUMAN (OTC)
 STRATTERA, ATOMOXETINE HYDROCHLORIDE
 SYMBYAX, FLUOXETINE HYDROCHLORIDE
 ZYPREXA ZYDIS, OLANZAPINE
 ZYPREXA, OLANZAPINE

**** F ****

FDC LTD

* FDC LTD
 CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE
 DORZOLAMIDE HYDROCHLORIDE, DORZOLAMIDE HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** F ******* FDC LTD**

LATANOPROST, LATANOPROST
 OFLOXACIN, OFLOXACIN
 OLOPATADINE HYDROCHLORIDE, OLOPATADINE HYDROCHLORIDE
 TIMOLOL MALEATE, TIMOLOL MALEATE

FDN CONSUMER

* 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

FERA PHARMS

* FERA PHARMACEUTICALS LLC
 TOBRAMYCIN, TOBRAMYCIN

FERA PHARMS LLC

* FERA PHARMACEUTICALS LLC
 DEXAMETHASONE, DEXAMETHASONE
 GENTAMICIN SULFATE, GENTAMICIN SULFATE

FERRER INTERNACIONAL

* FERRER INTERNACIONAL SA
 XEPI, OZENOXACIN

FERRING

* FERRING PHARMACEUTICALS INC
 ACTHREL, CORTICORELIN OVINE TRIFLUTATE
 CHORIONIC GONADOTROPIN, GONADOTROPIN, CHORIONIC
 ENDOMETRIN, PROGESTERONE
 FIRMAGON, DEGARELIX ACETATE
 MENOPUR, MENOTROPINS (FSH
 MINIRIN, DESMOPRESSIN ACETATE
 ZOMACTON, SOMATROPIN

FERRING PHARMS INC

* FERRING PHARMACEUTICALS INC
 CERVIDIL, DINOPROSTONE
 CLENPIQ, CITRIC ACID
 DDAVP (NEEDS NO REFRIGERATION), DESMOPRESSIN ACETATE
 DDAVP, DESMOPRESSIN ACETATE
 LYSTEDA, TRANEXAMIC ACID
 NOCDURNA, DESMOPRESSIN ACETATE
 STIMATE (NEEDS NO REFRIGERATION), DESMOPRESSIN ACETATE

FLAMINGO PHARMS

* FLAMINGO PHARMACEUTICALS LTD
 METRONIDAZOLE, METRONIDAZOLE
 PIROXICAM, PIROXICAM

FLEXION THERAPS INC

* FLEXION THERAPEUTICS INC
 ZILRETTA, TRIAMCINOLONE ACETONIDE

FOAMIX

* FOAMIX PHARMACEUTICALS INC
 AMZEEQ, MINOCYCLINE HYDROCHLORIDE

FOLDRX PHARMS

* FOLDRX PHARMACEUTICALS INC SUB PFIZER INC
 VYNDAMAX, TAFAMIDIS
 VYNDAQEL, TAFAMIDIS MEGLUMINE

FOREST LABS LLC

* FOREST LABORATORIES LLC
 NAMENDA XR, MEMANTINE HYDROCHLORIDE

FORTOVIA

* FORTOVIA THERAPEUTICS INC
 ORAVIG, MICONAZOLE
 SOLTAMOX, TAMOXIFEN CITRATE

FOSUN PHARMA

* FOSUN PHARMA USA INC

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** F ****

* FOSUN PHARMA USA INC
 DICYCLOMINE HYDROCHLORIDE, DICYCLOMINE HYDROCHLORIDE

FOUGERA PHARMS

* FOUGERA PHARMACEUTICALS INC
 ADAPALENE, ADAPALENE
 ALCLOMETASONE DIPROPIONATE, ALCLOMETASONE DIPROPIONATE
 AMCINONIDE, AMCINONIDE
 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
 CUTIVATE, FLUTICASONE PROPIONATE
 DESONIDE, DESONIDE
 DESOXIMETASONE, DESOXIMETASONE
 DIFLORASONE DIACETATE, DIFLORASONE DIACETATE
 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
 PANDEL, HYDROCORTISONE PROBUTATE
 PREDNICARBATE, PREDNICARBATE
 SOLARAZE, DICLOFENAC SODIUM
 SULFACETAMIDE SODIUM, SULFACETAMIDE SODIUM
 TERCONAZOLE, TERCONAZOLE
 TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE
 TYZINE, TETRAHYDROZOLINE HYDROCHLORIDE

FOUGERA PHARMS INC

* 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
 VEREGEN, SINECATECHINS

FRESENIUS

* FRESENIUS KABI DEUTSCHLAND GMBH
 INTRALIPID 10%, SOYBEAN OIL
 INTRALIPID 20%, SOYBEAN OIL
 INTRALIPID 30%, SOYBEAN OIL

* FRESENIUS KABI IPSUM SRL
 NAFCILLIN SODIUM, NAFCILLIN SODIUM

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** F ******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
ANASTROZOLE, ANASTROZOLE
ARGATROBAN, ARGATROBAN
ARSENIC TRIOXIDE, ARSENIC TRIOXIDE
ATROPINE SULFATE, ATROPINE SULFATE
AZITHROMYCIN, AZITHROMYCIN
AZTREONAM, AZTREONAM
BACITRACIN, BACITRACIN
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
CALCIUM GLUCONATE, CALCIUM GLUCONATE
CARBOPLATIN, CARBOPLATIN
CASPOFUNGIN ACETATE, CASPOFUNGIN ACETATE
CEFOTETAN, CEFOTETAN DISODIUM
CHLORAMPHENICOL SODIUM SUCCINATE, CHLORAMPHENICOL SODIUM SUCCINATE
CHLOROTHIAZIDE SODIUM, CHLOROTHIAZIDE SODIUM
CHORIONIC GONADOTROPIN, GONADOTROPIN, CHORIONIC
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
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% IN PLASTIC CONTAINER, DEXTROSE
DICYCLOMINE HYDROCHLORIDE, DICYCLOMINE HYDROCHLORIDE
DILAUDID, HYDROMORPHONE HYDROCHLORIDE
DIMENHYDRINATE, DIMENHYDRINATE
DIPHENHYDRAMINE HYDROCHLORIDE PRESERVATIVE FREE, DIPHENHYDRAMINE HYDROCHLORIDE
DIPRIVAN, PROPOFOL
DIPYRIDAMOLE, DIPYRIDAMOLE
DOXORUBICIN HYDROCHLORIDE, DOXORUBICIN HYDROCHLORIDE
DOXY 100, DOXYCYCLINE HYCLATE
DOXY 200, DOXYCYCLINE HYCLATE
EPIRUBICIN HYDROCHLORIDE, EPIRUBICIN HYDROCHLORIDE
ESMOLOL HYDROCHLORIDE, ESMOLOL HYDROCHLORIDE
ETOPOSIDE, ETOPOSIDE
FAMOTIDINE PRESERVATIVE FREE, FAMOTIDINE
FAMOTIDINE, FAMOTIDINE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** F ****

* FRESSENIUS KABI USA LLC
FENTANYL CITRATE PRESERVATIVE FREE, 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
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 IN PLASTIC CONTAINER, HEPARIN SODIUM
HEPARIN SODIUM PRESERVATIVE FREE, HEPARIN SODIUM
HEPARIN SODIUM, HEPARIN SODIUM
HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
IDARUBICIN HYDROCHLORIDE, IDARUBICIN HYDROCHLORIDE
IFOSFAMIDE, IFOSFAMIDE
INDOMETHACIN, INDOMETHACIN
IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE
KABIVEN IN PLASTIC CONTAINER, AMINO ACIDS
KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
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
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
MITOXANTRONE HYDROCHLORIDE, MITOXANTRONE HYDROCHLORIDE
MORPHINE SULFATE, MORPHINE SULFATE
MOXIFLOXACIN HYDROCHLORIDE, MOXIFLOXACIN HYDROCHLORIDE
NAROPIN, ROPIVACAINE HYDROCHLORIDE
NEBUPENT, PENTAMIDINE ISETHIONATE
NEOSTIGMINE METHYLSULFATE, NEOSTIGMINE METHYLSULFATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** F ****

* FRESENIUS KABI USA LLC
 NESACAINE, CHLOROPROCAINE HYDROCHLORIDE
 NESACAINE-MPF, CHLOROPROCAINE HYDROCHLORIDE
 OCTREOTIDE ACETATE (PRESERVATIVE FREE), OCTREOTIDE ACETATE
 OCTREOTIDE ACETATE, OCTREOTIDE ACETATE
 OMEGAVEN, FISH OIL TRIGLYCERIDES
 ONDANSETRON HYDROCHLORIDE PRESERVATIVE FREE, ONDANSETRON HYDROCHLORIDE
 ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
 OXALIPLATIN, OXALIPLATIN
 OXYTOCIN, OXYTOCIN
 PACLITAXEL, PACLITAXEL
 PALONOSETRON HYDROCHLORIDE, PALONOSETRON HYDROCHLORIDE
 PAMIDRONATE DISODIUM, PAMIDRONATE DISODIUM
 PENTAM, PENTAMIDINE ISETHIONATE
 PERIKABIVEN IN PLASTIC CONTAINER, AMINO ACIDS
 PHENYLEPHRINE HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE
 PIPERACILLIN AND TAZOBACTAM, PIPERACILLIN SODIUM
 POLOCAINE, MEPIVACAINE HYDROCHLORIDE
 POLOCAINE-MPF, MEPIVACAINE HYDROCHLORIDE
 POLYMYXIN B SULFATE, POLYMYXIN B SULFATE
 POTASSIUM CHLORIDE IN PLASTIC CONTAINER, POTASSIUM CHLORIDE
 POTASSIUM CHLORIDE, 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
 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 3% IN PLASTIC CONTAINER, SODIUM CHLORIDE
 STERILE WATER FOR INJECTION IN PLASTIC CONTAINER, STERILE WATER FOR INJECTION
 STERILE WATER FOR INJECTION, STERILE WATER FOR INJECTION
 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
 VIBISONE, CYANOCOBALAMIN
 VINBLASTINE SULFATE, VINBLASTINE SULFATE
 VINOURELBINE TARTRATE, VINOURELBINE TARTRATE
 XYLOCAINE W/ EPINEPHRINE, EPINEPHRINE
 XYLOCAINE, LIDOCAINE HYDROCHLORIDE
 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
 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

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

**** F ****

- * FRESENIUS MEDICAL CARE NORTH AMERICA
 DELFLEX W/ DEXTROSE 4.25% LOW MAGNESIUM LOW CALCIUM IN PLASTIC CONTAINER, CALCIUM
 PHOSLO GELCAPS, CALCIUM ACETATE
 PHOSLYRA, CALCIUM ACETATE
 SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, SODIUM CHLORIDE

FRONTIDA BIOPHARM

- * FRONTIDA BIOPHARM INC
 BISOPROLOL FUMARATE, BISOPROLOL FUMARATE
 CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE

**** G ****

G AND W LABS INC

- * G AND W LABORATORIES INC
 CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
 FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE

GALDERMA LABS

- * GALDERMA LABORATORIES INC
 CLOBEX, CLOBETASOL PROPIONATE
 EPIDUO FORTE, ADAPALENE

GALDERMA LABS LP

- * GALDERMA LABORATORIES L P
 CLOBEX, CLOBETASOL PROPIONATE
- * GALDERMA LABORATORIES LP
 CAPEX, FLUOCINOLONE ACETONIDE
 CLOBEX, CLOBETASOL PROPIONATE
 DESOWEN, DESONIDE
 DIFFERIN, ADAPALENE
 DIFFERIN, ADAPALENE (OTC)
 EPIDUO, ADAPALENE
 METROCREAM, METRONIDAZOLE
 METROGEL, METRONIDAZOLE
 METROLOTION, METRONIDAZOLE
 MIRVASO, BRIMONIDINE TARTRATE
 ORACEA, DOXYCYCLINE
 SOOLANTRA, IVERMECTIN
 TRI-LUMA, FLUOCINOLONE ACETONIDE
 VECTICAL, CALCITRIOL

GALDERMA R AND D

- * GALDERMA RESEARCH AND DEVELOPMENT INC
 AKLIEF, TRIFAROTENE

GALEN SPECIALTY

- * GALEN SPECIALTY PHARMA US LLC
 SYNERA, LIDOCAINE

GALT PHARMS

- * GALT PHARMACEUTICALS LLC
 DORAL, QUAZEPAM

GATE PHARMS

- * GATE PHARMACEUTICALS
 LINEZOLID, LINEZOLID

GD SEARLE

- * GD SEARLE LLC
 CELEBREX, CELECOXIB
 DAYPRO, OXAPROZIN

GD SEARLE LLC

- * GD SEARLE LLC
 ALDACTAZIDE, HYDROCHLOROTHIAZIDE
 ALDACTONE, SPIRONOLACTONE
 ARTHROTEC, DICLOFENAC SODIUM
 CYTOTEC, MISOPROSTOL
 FLAGYL, METRONIDAZOLE
 INSPRA, EPLERENONE
 LOMOTIL, ATROPINE SULFATE
 NORPACE CR, DISOPYRAMIDE PHOSPHATE
 NORPACE, DISOPYRAMIDE PHOSPHATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** G ****

* GD SEARLE LLC
SYNAREL, NAFARELIN ACETATE

GE HEALTHCARE

* GE HEALTHCARE
ADREVIEW, IOBENGUANE SULFATE I-123
CERETEC, TECHNETIUM TC-99M EXAMETAZIME KIT
CLARISCAN, GADOTERATE MEGLUMINE
INDIUM IN 111 OXYQUINOLINE, INDIUM IN-111 OXYQUINOLINE
MPI INDIUM DTPA IN 111, INDIUM IN-111 PENTETATE DISODIUM
MYOVUE 30ML, TECHNETIUM TC-99M TETROFOSMIN KIT
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
THALLOUS CHLORIDE TL 201, THALLOUS CHLORIDE TL-201
VISIPAQUE 270, IODIXANOL
VISIPAQUE 320, IODIXANOL
VIZAMYL, FLUTEMETAMOL F-18

GE HLTHCARE INC

* GE HEALTHCARE INC
DATSCAN, IOFLUPANE I-123

GENBIOPRO

* GENBIOPRO INC
MIFEPRISTONE, MIFEPRISTONE

GENENTECH

* GENENTECH INC
ERIVEDGE, VISMODEGIB
NUTROPIN AQ NUSPIN, SOMATROPIN

GENENTECH INC

* GENENTECH INC
COTELLIC, COBIMETINIB FUMARATE
ESBRIET, PIRFENIDONE
ROZLYTREK, ENTRECTINIB
XOFLUZA, BALOXAVIR MARBOXIL

GENERICIS

* GENERICIS INTERNATIONAL VENTURES ENTERPRISES LLC
DICYCLOMINE HYDROCHLORIDE, DICYCLOMINE HYDROCHLORIDE

GENEYORK PHARMS

* GENEYORK PHARMACEUTICALS GROUP LLC
PREDNISONE, PREDNISONE

GENUS

* GENUS LIFESCIENCES INC
HYCODAN, HOMATROPINE METHYLBROMIDE
HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
TIVORBEX, INDOMETHACIN

GENUS LIFESCIENCES

* GENUS LIFE SCIENCES INC
GOPRELTO, COCAINE HYDROCHLORIDE
OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
YOSPRALA, ASPIRIN

GENZYME

* GENZYME CORP
CEREZYME, IMIGLUCERASE
CLOLAR, CLOFARABINE
MOZOBIL, PLERIXAFOR
RENAGEL, SEVELAMER HYDROCHLORIDE
RENVELA, SEVELAMER CARBONATE
THYROGEN, THYROTROPIN ALFA

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** G ******GENZYME CORP**

* GENZYME CORP
 CAPRELSA, VANDETANIB
 CERDELGA, ELIGLUSTAT TARTRATE

GILEAD

* GILEAD SCIENCES INC
 CAYSTON, AZTREONAM
 EMTRIVA, EMTRICITABINE
 HEPSERA, ADEFOVIR DIPIVOXIL
 LETAIRIS, AMBRISENTAN
 RANEXA, RANOLAZINE
 TRUVADA, EMTRICITABINE

GILEAD SCIENCES

* GILEAD SCIENCES LLC
 ATRIPLA, EFAVIRENZ

GILEAD SCIENCES INC

* GILEAD SCIENCES INC
 BIKTARVY, BICTEGRAVIR SODIUM
 COMPLERA, EMTRICITABINE
 DESCOVY, EMTRICITABINE
 EPCLUSA, SOFOSBUVIR
 GENVOYA, COBICISTAT
 HARVONI, LEDIPASVIR
 ODEFSEY, EMTRICITABINE
 SOVALDI, SOFOSBUVIR
 STRIBILD, COBICISTAT
 TYBOST, COBICISTAT
 VEMLIDY, TENOFOVIR ALAFENAMIDE FUMARATE
 VIREAD, TENOFOVIR DISOPROXIL FUMARATE
 VOSEVI, SOFOSBUVIR
 ZYDELIG, IDELALISIB

GISKIT

* GISKIT BV
 EXEM FOAM KIT, AIR POLYMER-TYPE A

GLAND PHARMA LTD

* GLAND PHARMA LTD
 ADENOSINE, ADENOSINE
 AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE
 ARGATROBAN IN SODIUM CHLORIDE, ARGATROBAN
 AZELASTINE HYDROCHLORIDE, AZELASTINE HYDROCHLORIDE
 AZITHROMYCIN, AZITHROMYCIN
 BIMATOPROST, BIMATOPROST
 BROMFENAC SODIUM, BROMFENAC SODIUM
 CALCITRIOL, CALCITRIOL
 CARBOPLATIN, CARBOPLATIN
 CASPOFUNGIN ACETATE, CASPOFUNGIN ACETATE
 CISPLATIN, CISPLATIN
 CLOFARABINE, CLOFARABINE
 CYTARABINE, CYTARABINE
 DEFEROXAMINE MESYLATE, DEFEROXAMINE MESYLATE
 DEXRAZOXANE HYDROCHLORIDE, DEXRAZOXANE HYDROCHLORIDE
 DOXERCALCIFEROL, DOXERCALCIFEROL
 DOXORUBICIN HYDROCHLORIDE, DOXORUBICIN HYDROCHLORIDE
 ESMOLOL HYDROCHLORIDE, ESMOLOL HYDROCHLORIDE
 ETOMIDATE, ETOMIDATE
 FLUOROURACIL, FLUOROURACIL
 GEMCITABINE HYDROCHLORIDE, GEMCITABINE HYDROCHLORIDE
 GLYCOPYRROLATE, GLYCOPYRROLATE
 HALOPERIDOL DECANOATE, HALOPERIDOL DECANOATE
 HALOPERIDOL, HALOPERIDOL LACTATE
 HEPARIN SODIUM, HEPARIN SODIUM
 IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE
 KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
 LABETALOL HYDROCHLORIDE, LABETALOL HYDROCHLORIDE
 LEVETIRACETAM IN SODIUM CHLORIDE, LEVETIRACETAM

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** G ******* GLAND PHARMA LTD**

LEVOLEUCOVORIN CALCIUM, LEVOLEUCOVORIN CALCIUM
 MELPHALAN HYDROCHLORIDE, MELPHALAN HYDROCHLORIDE
 MEROPENEM, MEROPENEM
 MESNA, MESNA
 METHOCARBAMOL, METHOCARBAMOL
 METOPROLOL TARTRATE, METOPROLOL TARTRATE
 MIDAZOLAM HYDROCHLORIDE, MIDAZOLAM HYDROCHLORIDE
 NEOSTIGMINE METHYLSULFATE, NEOSTIGMINE METHYLSULFATE
 OLOPATADINE HYDROCHLORIDE, OLOPATADINE HYDROCHLORIDE
 ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
 OXALIPLATIN, OXALIPLATIN
 PACITAXEL, PACLITAXEL
 POLYMYXIN B SULFATE, POLYMYXIN B SULFATE
 ROCURONIUM BROMIDE, ROCURONIUM BROMIDE
 TEMSIROLIMUS, TEMSIROLIMUS
 TRANEXAMIC ACID, TRANEXAMIC ACID
 VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE
 VECURONIUM BROMIDE, VECURONIUM BROMIDE
 ZIPRASIDONE MESYLATE, ZIPRASIDONE MESYLATE
 ZOLEDRONIC ACID, ZOLEDRONIC ACID
 ZOLEDRONIC, ZOLEDRONIC ACID

GLASSHOUSE PHARMS*** GLASSHOUSE PHARMACEUTICALS LTD CANADA**

CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
 FLUOCINOLONE ACETONIDE, FLUOCINOLONE ACETONIDE
 FLUOCINONIDE, FLUOCINONIDE

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**

ABREVA, DOCOSANOL (OTC)
 AVODART, DUTASTERIDE
 BECONASE AQ, BECLOMETHASONE DIPROPIONATE MONOHYDRATE
 EPIVIR-HBV, LAMIVUDINE
 IMITREX STATDOSE, SUMATRIPTAN SUCCINATE
 IMITREX, SUMATRIPTAN
 IMITREX, SUMATRIPTAN SUCCINATE
 JALYN, DUTASTERIDE
 MALARONE PEDIATRIC, ATOVAQUONE
 MALARONE, ATOVAQUONE
 NICORETTE (MINT), NICOTINE POLACRILEX (OTC)
 NICORETTE, NICOTINE POLACRILEX (OTC)
 RELENZA, ZANAMIVIR
 VALTREX, VALACYCLOVIR HYDROCHLORIDE
 WELLBUTRIN SR, BUPROPION HYDROCHLORIDE

*** GLAXOSMITHKLINE CONSUMER HEALTHCARE HOLDINGS (US) LLC**

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 LIQUI-GELS, IBUPROFEN (OTC)

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

**** G ****

- * GLAXOSMITHKLINE CONSUMER HEALTHCARE HOLDINGS (US) LLC
 - 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)
 - ALAVERT, LORATADINE (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)
 - INFANT'S ADVIL, IBUPROFEN (OTC)
 - JUNIOR STRENGTH ADVIL, IBUPROFEN (OTC)
 - LAMISIL, TERBINAFFINE HYDROCHLORIDE (OTC)
 - LORATADINE, LORATADINE (OTC)
- * GLAXOSMITHKLINE INTELLECTUAL PROPERTY DEVELOPMENT LTD ENGLAND
 - ANORO ELLIPTA, UMECLIDINIUM BROMIDE
 - ARNUIITY ELLIPTA, FLUTICASONE FUROATE
 - KRINTAFEL, TAFENOQUINE SUCCINATE
 - TRELEGY ELLIPTA, FLUTICASONE FUROATE
- * GLAXOSMITHKLINE INTELLECTUAL PROPERTY LTD ENGLAND
 - SEREVENT, SALMETEROL XINAFOATE
 - VENTOLIN HFA, ALBUTEROL SULFATE
- * GLAXOSMITHKLINE LLC
 - ZEJULA, NIRAPARIB TOSYLATE

GLAXOSMITHKLINE CON

- * GLAXOSMITHKLINE CONSUMER HEALTH
 - TRANSDERM SCOP, SCOPOLAMINE

GLAXOSMITHKLINE CONS

- * GLAXOSMITHKLINE CONSUMER HEALTHCARE
 - ALLI, ORLISTAT (OTC)
 - EXCEDRIN (MIGRAINE), ACETAMINOPHEN (OTC)
 - FLONASE ALLERGY RELIEF, FLUTICASONE PROPIONATE (OTC)
 - FLONASE SENSIMIST ALLERGY RELIEF, FLUTICASONE FUROATE (OTC)
 - LAMISIL AT, TERBINAFFINE (OTC)
 - LAMISIL AT, TERBINAFFINE HYDROCHLORIDE (OTC)
 - NICORETTE, NICOTINE POLACRILEX (OTC)
 - VOLTAREN, DICLOFENAC SODIUM

GLAXOSMITHKLINE LLC

- * GLAXOSMITHKLINE LLC
 - AMERGE, NARATRIPTAN HYDROCHLORIDE
 - DYAZIDE, HYDROCHLOROTHIAZIDE
 - FLOLAN, EPOPROSTENOL SODIUM
 - LAMICTAL CD, LAMOTRIGINE
 - LAMICTAL ODT, LAMOTRIGINE
 - LAMICTAL XR, LAMOTRIGINE
 - LAMICTAL, LAMOTRIGINE
 - MEPRON, ATOVAQUONE
 - REQUIP XL, ROPINIROLE HYDROCHLORIDE
 - REQUIP, ROPINIROLE HYDROCHLORIDE
 - RYTHMOL SR, PROPAFENONE HYDROCHLORIDE

GLENMARK

- * GLENMARK THERAPEUTICS INC USA
 - ECOZA, ECONAZOLE NITRATE

GLENMARK GENERICS

- * GLENMARK GENERICS INC USA
 - ADAPALENE, ADAPALENE
 - ADAPALENE, ADAPALENE (OTC)
 - BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
 - IMIQUIMOD, IMIQUIMOD
 - MOMETASONE FUROATE, MOMETASONE FUROATE
 - NIZATIDINE, NIZATIDINE
 - ZONISAMIDE, ZONISAMIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** G ****

- * GLENMARK GENERICS LIMITED
 - BRIELLYN, ETHINYL ESTRADIOL
 - FLUTICASONE PROPIONATE, FLUTICASONE PROPIONATE
 - LEVOCETIRIZINE DIHYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE
- * GLENMARK GENERICS LTD
 - ACAMPROSATE CALCIUM, ACAMPROSATE CALCIUM
 - ALCLOMETASONE DIPROPIONATE, ALCLOMETASONE DIPROPIONATE
 - ALYACEN 1/35, ETHINYL ESTRADIOL
 - ALYACEN 7/7/7, ETHINYL ESTRADIOL
 - ASHLYNA, ETHINYL ESTRADIOL
 - ATOVAQUONE AND PROGUANIL HYDROCHLORIDE, ATOVAQUONE
 - CARVEDILOL, CARVEDILOL
 - CICLOPIROX, CICLOPIROX
 - CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE
 - DESOXIMETASONE, DESOXIMETASONE
 - ESZOPICLONE, ESZOPICLONE
 - FELODIPINE, FELODIPINE
 - FLUCONAZOLE, FLUCONAZOLE
 - FLUOCINONIDE, FLUOCINONIDE
 - HEATHER, NORETHINDRONE
 - HYDROCORTISONE BUTYRATE, HYDROCORTISONE BUTYRATE
 - LAMOTRIGINE, LAMOTRIGINE
 - LEVOFLOXACIN, LEVOFLOXACIN
 - LEVONORGESTREL AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
 - LITHIUM CARBONATE, LITHIUM CARBONATE
 - MARLISSA, ETHINYL ESTRADIOL
 - MELOXICAM, MELOXICAM
 - METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 - MOEXIPRIL HYDROCHLORIDE, MOEXIPRIL HYDROCHLORIDE
 - MOMETASONE FUROATE, MOMETASONE FUROATE
 - MONTELUKAST SODIUM, MONTELUKAST SODIUM
 - NAPROXEN, NAPROXEN
 - NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
 - NORETHINDRONE, NORETHINDRONE
 - NORGESTIMATE AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
 - OMEPRAZOLE, OMEPRAZOLE
 - ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
 - ONDANSETRON, ONDANSETRON
 - PRAVASTATIN SODIUM, PRAVASTATIN SODIUM
 - RIZATRIPTAN BENZOATE, RIZATRIPTAN BENZOATE
 - ROPINIROLE HYDROCHLORIDE, ROPINIROLE HYDROCHLORIDE
 - SULFAMETHOXAZOLE AND TRIMETHOPRIM, SULFAMETHOXAZOLE
 - TERBINAFINE HYDROCHLORIDE, TERBINAFINE HYDROCHLORIDE
 - THEOPHYLLINE, THEOPHYLLINE
 - TOPIRAMATE, TOPIRAMATE
 - TRANDOLAPRIL AND VERAPAMIL HYDROCHLORIDE, TRANDOLAPRIL
 - TROSPIUM CHLORIDE, TROSPIUM CHLORIDE
 - URSODIOL, URSODIOL
 - VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE
 - VIORELE, DESOGESTREL
 - ZOLMITRIPTAN, ZOLMITRIPTAN
- * GLENMARK GENERICS LTD INDIA
 - INDOMETHACIN, INDOMETHACIN
 - NORETHINDRONE ACETATE, NORETHINDRONE ACETATE
 - PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE
- GLENMARK PHARMS**
- * GLENMARK PHARMACEUTICALS INC
 - TERIFLUNOMIDE, TERIFLUNOMIDE
- * GLENMARK PHARMACEUTICALS INC USA
 - CICLOPIROX, CICLOPIROX
 - CLOTRIMAZOLE, CLOTRIMAZOLE
 - MUPIROCIN, MUPIROCIN
- * GLENMARK PHARMACEUTICALS LTD
 - MOEXIPRIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
- * GLENMARK PHARMACEUTICALS SA

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** G ****

* GLENMARK PHARMACEUTICALS SA

ABIRATERONE ACETATE, ABIRATERONE ACETATE
 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
 DESONIDE, DESONIDE
 ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM
 HAILEY 1.5/30, ETHINYL ESTRADIOL
 HAILEY FE 1.5/30, ETHINYL ESTRADIOL
 LINEZOLID, LINEZOLID
 NORGESTIMATE AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
 ROSUVASTATIN CALCIUM, ROSUVASTATIN CALCIUM
 TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE

GLENMARK PHARMS INC

* GLENMARK PHARMACEUTICALS INC USA

CALCIPOTRIENE, CALCIPOTRIENE
 FULVESTRANT, FULVESTRANT
 LITHIUM CARBONATE, LITHIUM CARBONATE
 MUPIROCIN, MUPIROCIN CALCIUM
 RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
 SOLIFENACIN SUCCINATE, SOLIFENACIN SUCCINATE

GLENMARK PHARMS LTD

* GLENMARK PHARMACEUTICALS LTD

ADAPALENE AND BENZOYL PEROXIDE, ADAPALENE
 AMLODIPINE AND OLMESARTAN MEDOXOMIL, AMLODIPINE BESYLATE
 ATOMOXETINE HYDROCHLORIDE, ATOMOXETINE HYDROCHLORIDE
 CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
 COLESEVELAM HYDROCHLORIDE, COLESEVELAM HYDROCHLORIDE
 DESMOPRESSIN ACETATE, DESMOPRESSIN ACETATE
 DESONIDE, DESONIDE
 DICLOFENAC SODIUM, DICLOFENAC SODIUM
 DROSPIRENONE AND ETHINYL ESTRADIOL, DROSPIRENONE
 ESTRADIOL, ESTRADIOL
 EZETIMIBE AND SIMVASTATIN, EZETIMIBE
 EZETIMIBE, EZETIMIBE
 FENOFIBRATE (MICRONIZED), FENOFIBRATE
 FLUCINOLONE ACETONIDE, FLUCINOLONE ACETONIDE
 FLUCINONIDE ACETONIDE, FLUCINOLONE ACETONIDE
 FROVATRIPTAN SUCCINATE, FROVATRIPTAN SUCCINATE
 GABAPENTIN, GABAPENTIN
 HAILEY FE 1/20, ETHINYL ESTRADIOL
 HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
 HYDROCORTISONE VALERATE, HYDROCORTISONE VALERATE
 INDOMETHACIN, INDOMETHACIN
 LAMOTRIGINE, LAMOTRIGINE
 LEVONORGESTREL AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
 LEVONORGESTREL, LEVONORGESTREL (OTC)
 LIDOCAINE, LIDOCAINE
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 NAPROXEN SODIUM, NAPROXEN SODIUM
 NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL AND FERROUS FUMARATE, ETHINYL ESTRADIOL
 NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
 NYSTATIN AND TRIAMCINOLONE ACETONIDE, NYSTATIN
 OLMESARTAN MEDOXOMIL, OLMESARTAN MEDOXOMIL
 OXCARBAZEPINE, OXCARBAZEPINE
 PIMECROLIMUS, PIMECROLIMUS
 POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
 PROPAFENONE HYDROCHLORIDE, PROPAFENONE HYDROCHLORIDE
 RALOXIFENE HYDROCHLORIDE, RALOXIFENE HYDROCHLORIDE
 RANOLAZINE, RANOLAZINE
 RILUZOLE, RILUZOLE
 RIZATRIPTAN BENZOATE, RIZATRIPTAN BENZOATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** G ****

* GLENMARK PHARMACEUTICALS LTD
 RUFINAMIDE, RUFINAMIDE
 SEVELAMER HYDROCHLORIDE, SEVELAMER HYDROCHLORIDE
 TACROLIMUS, TACROLIMUS
 TELMISARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 TELMISARTAN, TELMISARTAN
 TRETINOIN, TRETINOIN
 TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE
 VORICONAZOLE, VORICONAZOLE

GLENMARK PHARMS SA

* GLENMARK PHARMACEUTICALS SA SWITZERLAND
 ACYCLOVIR, ACYCLOVIR
 APREPITANT, APREPITANT
 ASPIRIN AND DIPYRIDAMOLE, ASPIRIN
 NITROGLYCERIN, NITROGLYCERIN

GLOBAL BLOOD THERAPYS

* GLOBAL BLOOD THERAPEUTICS INC
 OXBRYTA, VOXELOTOR

GRANULES INDIA

* GRANULES INDIA LTD
 IBUPROFEN, IBUPROFEN (OTC)
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 NAPROXEN SODIUM, NAPROXEN SODIUM (OTC)

GRANULES INDIA LTD

* GRANULES INDIA LTD
 ACETAMINOPHEN, ACETAMINOPHEN (OTC)
 CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 FEXOFENADINE HYDROCHLORIDE ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
 GABAPENTIN, GABAPENTIN
 IBUPROFEN, IBUPROFEN
 IBUPROFEN, IBUPROFEN (OTC)
 LORATADINE, LORATADINE (OTC)
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 METHOCARBAMOL, METHOCARBAMOL
 RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE (OTC)

GRANULES PHARMS

* GRANULES PHARMACEUTICALS INC
 AMPHETAMINE SULFATE, AMPHETAMINE SULFATE
 BUTALBITAL AND ACETAMINOPHEN, ACETAMINOPHEN
 COLCHICINE, COLCHICINE
 HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
 METHYLERGONOVINE MALEATE, METHYLERGONOVINE MALEATE
 METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
 OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN
 POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
 VALGANCICLOVIR HYDROCHLORIDE, VALGANCICLOVIR HYDROCHLORIDE

GRAVITI PHARMS

* GRAVITI PHARMACEUTICALS PRIVATE LTD
 ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM
 BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
 ESCITALOPRAM OXALATE, ESCITALOPRAM OXALATE
 FENOFIBRATE, FENOFIBRATE
 FENOFIBRIC ACID, CHOLINE FENOFIBRATE

GUARDIAN DRUG

* GUARDIAN DRUG CO
 GUAIFENESIN, GUAIFENESIN (OTC)
 IBUPROFEN, IBUPROFEN (OTC)
 LORATADINE, LORATADINE (OTC)

GUERBET

* GUERBET LLC
 DOTAREM, GADOTERATE MEGLUMINE
 LIPIODOL, ETHIODIZED OIL

GW RES LTD

* GW RESEARCH LTD
 EPIDIOLEX, CANNABIDIOL

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

**** G ****

HANFORD GC

- * GC HANFORD MANUFACTURING CO
 AMPICILLIN AND SULBACTAM, AMPICILLIN SODIUM
 AMPICILLIN SODIUM, AMPICILLIN SODIUM
 PENICILLIN G POTASSIUM, PENICILLIN G POTASSIUM

POHL BOSKAMP

- * G POHL BOSKAMP GMBH AND CO KG
 GONITRO, NITROGLYCERIN

**** H ****

HAEMONETICS

- * HAEMONETICS CORP
 SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, SODIUM CHLORIDE

HAINAN POLY PHARM

- * HAINAN POLY PHARMACEUTICAL CO LTD
 AZITHROMYCIN, AZITHROMYCIN
 EPTIFIBATIDE, EPTIFIBATIDE
 GANCICLOVIR SODIUM, GANCICLOVIR SODIUM
 LEVETIRACETAM, LEVETIRACETAM
 VORICONAZOLE, VORICONAZOLE

HALOCARBON PRODS

- * HALOCARBON PRODUCTS CORP
 ISOFLURANE, ISOFLURANE
 SEVOFLURANE, SEVOFLURANE

HALOZYME THERAP

- * HALOZYME THERAPEUTICS INC
 HYLENEX RECOMBINANT, HYALURONIDASE RECOMBINANT HUMAN

HAMELN PHARMA PLUS

- * HAMELN PHARMA PLUS GMBH
 PENTETATE CALCIUM TRISODIUM, PENTETATE CALCIUM TRISODIUM
 PENTETATE ZINC TRISODIUM, PENTETATE ZINC TRISODIUM

HANGZHOU BINJIANG

- * HANGZHOU MINSHENG BINJIANG PHARMACEUTICAL CO LTD
 ALENDRONATE SODIUM, ALENDRONATE SODIUM

HANSAMED INC

- * HANSAMED INC
 ULTACAN, ARTICAINE HYDROCHLORIDE

HARMONY

- * HARMONY BIOSCIENCES LLC
 WAKIX, PITOLISANT HYDROCHLORIDE

HARRIS PHARM

- * HARRIS PHARMACEUTICAL INC
 FLUCONAZOLE, FLUCONAZOLE
 TERBINAFINE HYDROCHLORIDE, TERBINAFINE HYDROCHLORIDE

HBT LABS INC

- * HBT LABS INC
 FULVESTRANT, FULVESTRANT

HEBEI CHANGSHAN

- * HEBEI CHANGSHAN BIOCHEMICAL PHARMACEUTICAL CO LTD
 AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
 SILDENAFIL CITRATE, SILDENAFIL CITRATE

HEC PHARM

- * HEC PHARM USA INC
 CLARITHROMYCIN, CLARITHROMYCIN
 LEVOFLOXACIN, LEVOFLOXACIN
 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

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** H ****

* HELSINN HEALTHCARE SA
AKYNZEO, FOSNETUPITANT CHLORIDE HYDROCHLORIDE
AKYNZEO, NETUPITANT
ALOXI, PALONOSETRON HYDROCHLORIDE

HERCON PHARM

* HERCON PHARMACEUTICAL LLC
NITROGLYCERIN, NITROGLYCERIN

HERITAGE

* HERITAGE PHARMACEUTICALS INC DBA AVET PHARMACEUTICALS INC
INDOMETHACIN, INDOMETHACIN

HERITAGE LIFE

* HERITAGE LIFE SCIENCES BARBADOS INC
CLOZARIL, CLOZAPINE

HERITAGE PHARMA

* HERITAGE PHARMA LABS INC
ACETAMINOPHEN, ACETAMINOPHEN (OTC)
ACETAZOLAMIDE, ACETAZOLAMIDE
AMANTADINE HYDROCHLORIDE, AMANTADINE HYDROCHLORIDE
BETHANECHOL CHLORIDE, BETHANECHOL CHLORIDE
BUMETANIDE, BUMETANIDE
BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
BUSPIRONE HYDROCHLORIDE, BUSPIRONE HYDROCHLORIDE
CYPROHEPTADINE HYDROCHLORIDE, CYPROHEPTADINE HYDROCHLORIDE
DESMOPRESSIN ACETATE, DESMOPRESSIN ACETATE
DIFLUNISAL, DIFLUNISAL
DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
DOXAZOSIN MESYLATE, DOXAZOSIN MESYLATE
DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
ENALAPRIL MALEATE, ENALAPRIL MALEATE
HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE
HYDROXYZINE PAMOATE, HYDROXYZINE PAMOATE
INDOMETHACIN, INDOMETHACIN
LABETALOL HYDROCHLORIDE, LABETALOL HYDROCHLORIDE
LITHIUM CARBONATE, LITHIUM CARBONATE
METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
METHIMAZOLE, METHIMAZOLE
METHYLDOPA, METHYLDOPA
NADOLOL, NADOLOL
NIFEDIPINE, NIFEDIPINE
PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE
RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
RISPERIDONE, RISPERIDONE
TERBINAFINE HYDROCHLORIDE, TERBINAFINE HYDROCHLORIDE
THEOPHYLLINE, THEOPHYLLINE
VIVACTIL, PROTRIPTYLINE HYDROCHLORIDE

HERITAGE PHARMS INC

* HERITAGE PHARMACEUTICALS INC
ACETAZOLAMIDE, ACETAZOLAMIDE
ACYCLOVIR, ACYCLOVIR
CALCIUM ACETATE, CALCIUM ACETATE
CARISOPRODOL AND ASPIRIN, ASPIRIN
CLINDAMYCIN PALMITATE HYDROCHLORIDE, CLINDAMYCIN PALMITATE HYDROCHLORIDE
DESIPRAMINE HYDROCHLORIDE, DESIPRAMINE HYDROCHLORIDE
DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
DOXYCYCLINE, DOXYCYCLINE
DUTASTERIDE, DUTASTERIDE
ETHOSUXIMIDE, ETHOSUXIMIDE
FELODIPINE, FELODIPINE
GLIPIZIDE AND METFORMIN HYDROCHLORIDE, GLIPIZIDE
GLYBURIDE, GLYBURIDE
GLYCOPYRROLATE, GLYCOPYRROLATE
HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
KETOPROFEN, KETOPROFEN
LEFLUNOMIDE, LEFLUNOMIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** H ****

* HERITAGE PHARMACEUTICALS INC
 METRONIDAZOLE, METRONIDAZOLE
 MOEXIPRIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 NARATRIPTAN, NARATRIPTAN HYDROCHLORIDE
 NIMODIPINE, NIMODIPINE
 NYSTATIN, NYSTATIN
 PAROMOMYCIN SULFATE, PAROMOMYCIN SULFATE
 TROSPIMUM CHLORIDE, TROSPIMUM CHLORIDE
 VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE
 VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE

HERON THERAPS INC

* HERON THERAPEUTICS INC
 CINVANTI, APREPITANT
 SUSTOL, GRANISETRON

HETERO LABS LTD

* HETERO LABS LTD
 DROSPIRENONE AND ETHINYL ESTRADIOL, DROSPIRENONE

HETERO LABS LTD III

* HETERO LABS LTD UNIT III
 ABACAVIR SULFATE, ABACAVIR SULFATE
 ATOVAQUONE, ATOVAQUONE
 CHILDREN'S CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE
 CLOBAZAM, CLOBAZAM
 DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE
 EFAVIRENZ, EFAVIRENZ
 ESCITALOPRAM OXALATE, ESCITALOPRAM OXALATE
 ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM
 FENOFIBRATE, FENOFIBRATE
 FINASTERIDE, FINASTERIDE
 HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
 HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE
 INDOMETHACIN, INDOMETHACIN
 LAMIVUDINE AND ZIDOVUDINE, LAMIVUDINE
 LEVETIRACETAM, LEVETIRACETAM
 LEVOCETIRIZINE DIHYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE
 LEVOCETIRIZINE HYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE
 LITHIUM CARBONATE, LITHIUM CARBONATE
 METHOCARBAMOL, METHOCARBAMOL
 NEVIRAPINE, NEVIRAPINE
 OMEPRAZOLE, OMEPRAZOLE
 OSELTAMIVIR PHOSPHATE, OSELTAMIVIR PHOSPHATE
 PREGABALIN, PREGABALIN
 RITONAVIR, RITONAVIR
 ROFLUMILAST, ROFLUMILAST
 SIMVASTATIN, SIMVASTATIN
 TADALAFIL, TADALAFIL
 TENOFOVIR DISOPROXIL FUMARATE, TENOFOVIR DISOPROXIL FUMARATE
 TOLTERODINE TARTRATE, TOLTERODINE TARTRATE
 TORSEMIDE, TORSEMIDE
 ZIDOVUDINE, ZIDOVUDINE

HETERO LABS LTD V

* HETERO LABS LTD UNIT V
 ACYCLOVIR, ACYCLOVIR
 ARIPIPRAZOLE, ARIPIPRAZOLE
 CHILDREN'S FEXOFENADINE HYDROCHLORIDE ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
 DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
 ENTECAVIR, ENTECAVIR
 FAMCICLOVIR, FAMCICLOVIR
 FEXOFENADINE HYDROCHLORIDE ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
 IRBESARTAN, IRBESARTAN
 LAMIVUDINE AND ZIDOVUDINE, LAMIVUDINE
 LAMIVUDINE, LAMIVUDINE
 LEVOFLOXACIN, LEVOFLOXACIN
 LINEZOLID, LINEZOLID

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** H ****

* HETERO LABS LTD UNIT V
 LOSARTAN POTASSIUM, LOSARTAN POTASSIUM
 MONTELUKAST SODIUM, MONTELUKAST SODIUM
 PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
 ROSUVASTATIN CALCIUM, ROSUVASTATIN CALCIUM
 SILDENAFIL CITRATE, SILDENAFIL CITRATE
 SILODOSIN, SILODOSIN
 TELMISARTAN, TELMISARTAN
 TETRABENAZINE, TETRABENAZINE
 VALACYCLOVIR HYDROCHLORIDE, VALACYCLOVIR HYDROCHLORIDE
 VALGANCICLOVIR HYDROCHLORIDE, VALGANCICLOVIR HYDROCHLORIDE
 VALSARTAN, VALSARTAN

HEYL CHEMISCH

* HEYL CHEMISCH PHARMAZEUTISCHE FABRIK
 RADIOGARDASE (PRUSSIAN BLUE), FERRIC HEXACYANOFERRATE (II)

HI TECH

* HI-TECH PHARMACAL CO INC
 BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
 BIMATOPROST, BIMATOPROST
 BROMFENAC SODIUM, BROMFENAC SODIUM
 CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
 DESONIDE, DESONIDE
 DICLOFENAC SODIUM, DICLOFENAC SODIUM
 IBUPROFEN, IBUPROFEN
 MEGESTROL ACETATE, MEGESTROL ACETATE
 MORPHINE SULFATE, MORPHINE SULFATE
 OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
 PHENYLEPHRINE HYDROCHLORIDE AND PROMETHAZINE HYDROCHLORIDE, PHENYLEPHRINE
 TRIFLURIDINE, TRIFLURIDINE

* HI-TECH PHARMACAL CO INC AN AKORN CO
 AZELASTINE HYDROCHLORIDE, AZELASTINE HYDROCHLORIDE
 FLUTICASONE PROPIONATE, FLUTICASONE PROPIONATE (OTC)
 LOTEPIREDNOL ETABONATE, LOTEPIREDNOL ETABONATE

HI TECH PHARMA

* HI TECH PHARMACAL CO INC
 ACETAMINOPHEN AND CODEINE PHOSPHATE, ACETAMINOPHEN
 ACYCLOVIR, ACYCLOVIR
 ALBUTEROL SULFATE, ALBUTEROL SULFATE
 AMANTADINE HYDROCHLORIDE, AMANTADINE HYDROCHLORIDE
 CALCIPOTRIENE, CALCIPOTRIENE
 CHLORHEXIDINE GLUCONATE, CHLORHEXIDINE GLUCONATE
 CICLOPIROX, CICLOPIROX
 CIMETIDINE HYDROCHLORIDE, CIMETIDINE HYDROCHLORIDE
 CORMAX, CLOBETASOL PROPIONATE
 DORZOLAMIDE HYDROCHLORIDE AND TIMOLOL MALEATE, DORZOLAMIDE HYDROCHLORIDE
 DORZOLAMIDE HYDROCHLORIDE, DORZOLAMIDE HYDROCHLORIDE
 EMBELINE E, CLOBETASOL PROPIONATE
 EMBELINE, CLOBETASOL PROPIONATE
 FLUTICASONE PROPIONATE, FLUTICASONE PROPIONATE
 GABAPENTIN, GABAPENTIN
 HYDROCODONE BITARTRATE AND HOMATROPINE METHYLBROMIDE, HOMATROPINE METHYLBROMIDE
 HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE
 LACTULOSE, LACTULOSE
 LEVOCARNITINE, LEVOCARNITINE
 LEVOFLOXACIN, LEVOFLOXACIN
 LIDOCAINE AND PRILOCAINE, LIDOCAINE
 LIDOCAINE HYDROCHLORIDE, LIDOCAINE HYDROCHLORIDE
 LOPERAMIDE HYDROCHLORIDE, LOPERAMIDE HYDROCHLORIDE (OTC)
 MIDAZOLAM HYDROCHLORIDE, MIDAZOLAM HYDROCHLORIDE
 MINOXIDIL (FOR MEN), MINOXIDIL (OTC)
 MINOXIDIL (FOR WOMEN), MINOXIDIL (OTC)
 NYSTATIN, NYSTATIN
 OFLOXACIN, OFLOXACIN
 PREDNISOLONE SODIUM PHOSPHATE, PREDNISOLONE SODIUM PHOSPHATE
 PROMETHAZINE HYDROCHLORIDE AND CODEINE PHOSPHATE, CODEINE PHOSPHATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** H ****

- * HI TECH PHARMACAL CO INC
 PROMETHAZINE HYDROCHLORIDE AND DEXTROMETHORPHAN HYDROBROMIDE, DEXTROMETHORPHAN
 PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
 RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
 SULFAMETHOXAZOLE AND TRIMETHOPRIM, SULFAMETHOXAZOLE
 TIMOLOL MALEATE, TIMOLOL MALEATE
 VOSOL HC, ACETIC ACID, GLACIAL
 VOSOL, ACETIC ACID, GLACIAL
- HI TECH PHARMA CO**
- * HI TECH PHARMACAL CO INC
 FLUNISOLIDE, FLUNISOLIDE
 PREDNISOLONE, PREDNISOLONE
- HI-TECH PHARMA CO**
- * HI-TECH PHARMACAL CO INC
 FAMOTIDINE, FAMOTIDINE
 GATIFLOXACIN, GATIFLOXACIN
 LORAZEPAM, LORAZEPAM
 PROMETHAZINE HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE W/CODEINE PHOSPHATE, CODEINE
- HI-TECH PHARMACAL**
- * HI-TECH PHARMACAL CO INC
 LEVETIRACETAM, LEVETIRACETAM
- HIGH TECH PHARMA**
- * HIGH TECHNOLOGY PHARMACAL CO INC
 VALPROIC ACID, VALPROIC ACID
- HIKMA**
- * HIKMA FARMACEUTICA LDA
 CEFOTAXIME, CEFOTAXIME SODIUM
- * HIKMA FARMACEUTICA PORTUGAL SA
 CEFOTETAN, CEFOTETAN DISODIUM
 CEFTRIAZONE SODIUM, CEFTRIAZONE SODIUM
 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
 CEFACLOR, CEFACLOR
 CEFADROXIL, CEFADROXIL/CEFADROXIL HEMIHYDRATE
 CEPHALEXIN, CEPHALEXIN
 CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE
 GLYBURIDE (MICRONIZED), GLYBURIDE
- * HIKMA PHARMACEUTICALS INTERNATIONAL LTD
 ATIVAN, LORAZEPAM
 CAFECIT, CAFFEINE CITRATE
 CIMETIDINE, CIMETIDINE
 CODEINE SULFATE, CODEINE SULFATE
 DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE
 DIGOXIN, DIGOXIN
 DOPRAM, DOXAPRAM HYDROCHLORIDE
 FUROSEMIDE, FUROSEMIDE
 HEPARIN SODIUM, HEPARIN SODIUM
 MORPHINE SULFATE, MORPHINE SULFATE
 ROBAXIN, METHOCARBAMOL
 RUFINAMIDE, RUFINAMIDE
 TERAZOSIN HYDROCHLORIDE, TERAZOSIN HYDROCHLORIDE
- * HIKMA PHARMACEUTICALS LLC
 PALONOSETRON HYDROCHLORIDE, PALONOSETRON HYDROCHLORIDE
 PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
- * HIKMA PHARMACEUTICALS USA INC
 ACARBOSE, ACARBOSE
 ALENDRONATE SODIUM, ALENDRONATE SODIUM
 ALOSETRON HYDROCHLORIDE, ALOSETRON HYDROCHLORIDE
 ALPRAZOLAM, ALPRAZOLAM

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** H ****

* HIKMA PHARMACEUTICALS USA INC
 ANASTROZOLE, ANASTROZOLE
 AZELASTINE HYDROCHLORIDE, AZELASTINE HYDROCHLORIDE
 BALSALAZIDE DISODIUM, BALSALAZIDE DISODIUM
 BOSENTAN, BOSENTAN
 BUPRENORPHINE HYDROCHLORIDE AND NALOXONE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE
 BUPRENORPHINE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE
 BUTORPHANOL TARTRATE, BUTORPHANOL TARTRATE
 CALCITRIOL, CALCITRIOL
 CALCIUM ACETATE, CALCIUM ACETATE
 CAPECITABINE, CAPECITABINE
 CEVIMELINE HYDROCHLORIDE, CEVIMELINE HYDROCHLORIDE
 CILOSTAZOL, CILOSTAZOL
 CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE
 CLARITHROMYCIN, CLARITHROMYCIN
 CLOBAZAM, CLOBAZAM
 CLOTRIMAZOLE, CLOTRIMAZOLE
 CYCLOPHOSPHAMIDE, CYCLOPHOSPHAMIDE
 DALFAMPRIDINE, DALFAMPRIDINE
 DESVENLAFAXINE SUCCINATE, DESVENLAFAXINE SUCCINATE
 DEXAMETHASONE INTENSOL, DEXAMETHASONE
 DEXAMETHASONE, DEXAMETHASONE
 DIAZEPAM INTENSOL, DIAZEPAM
 DIAZEPAM, DIAZEPAM
 DIPHENOXYLATE HYDROCHLORIDE AND ATROPINE SULFATE, ATROPINE SULFATE
 DOLOPHINE HYDROCHLORIDE, METHADONE HYDROCHLORIDE
 DOXERCALCIFEROL, DOXERCALCIFEROL
 DURAMORPH PF, MORPHINE SULFATE
 ETHACRYNIC ACID, ETHACRYNIC ACID
 EVEROLIMUS, EVEROLIMUS
 EXEMESTANE, EXEMESTANE
 FAMCICLOVIR, FAMCICLOVIR
 FEBUXOSTAT, FEBUXOSTAT
 FENTANYL CITRATE PRESERVATIVE FREE, FENTANYL CITRATE
 FLECAINIDE ACETATE, FLECAINIDE ACETATE
 FLUCYTOSINE, FLUCYTOSINE
 FLUTICASONE PROPIONATE, FLUTICASONE PROPIONATE
 FLUTICASONE PROPIONATE, FLUTICASONE PROPIONATE (OTC)
 FUROSEMIDE, FUROSEMIDE
 GALANTAMINE HYDROBROMIDE, GALANTAMINE HYDROBROMIDE
 GRANISETRON HYDROCHLORIDE, GRANISETRON HYDROCHLORIDE
 HYDROMORPHONE HYDROCHLORIDE, HYDROMORPHONE HYDROCHLORIDE
 IMATINIB MESYLATE, IMATINIB MESYLATE
 IMIPRAMINE PAMOATE, IMIPRAMINE PAMOATE
 INFUMORPH, MORPHINE SULFATE
 IPRATROPIUM BROMIDE, IPRATROPIUM BROMIDE
 IRBESARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 LACTULOSE, LACTULOSE
 LETROZOLE, LETROZOLE
 LEUCOVORIN CALCIUM, LEUCOVORIN CALCIUM
 LIDOCAINE HYDROCHLORIDE, LIDOCAINE HYDROCHLORIDE
 LIDOCAINE VISCOUS, LIDOCAINE HYDROCHLORIDE
 LINEZOLID, LINEZOLID
 LITHIUM CARBONATE, LITHIUM CARBONATE
 LITHIUM CITRATE, LITHIUM CITRATE
 LORAZEPAM INTENSOL, LORAZEPAM
 LOSARTAN POTASSIUM AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 MEGESTROL ACETATE, MEGESTROL ACETATE
 MEPERIDINE HYDROCHLORIDE, MEPERIDINE HYDROCHLORIDE
 MERCAPTOPYRINE, MERCAPTOPYRINE
 METHADONE HYDROCHLORIDE INTENSOL, METHADONE HYDROCHLORIDE
 METHADONE HYDROCHLORIDE, METHADONE HYDROCHLORIDE
 METHAMPHETAMINE HYDROCHLORIDE, METHAMPHETAMINE HYDROCHLORIDE
 METHOTREXATE SODIUM, METHOTREXATE SODIUM
 MIDAZOLAM HYDROCHLORIDE, MIDAZOLAM HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** H ****

* HIKMA PHARMACEUTICALS USA INC
 MONTELUKAST SODIUM, MONTELUKAST SODIUM
 MORPHINE SULFATE, MORPHINE SULFATE
 MYCOPHENOLATE MOFETIL, MYCOPHENOLATE MOFETIL
 NAPROXEN, NAPROXEN
 NARATRIPTAN, NARATRIPTAN HYDROCHLORIDE
 ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
 OXCARBAZEPINE, OXCARBAZEPINE
 OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
 OXYMORPHONE HYDROCHLORIDE, OXYMORPHONE HYDROCHLORIDE
 PERINDOPRIL ERBUMINE, PERINDOPRIL ERBUMINE
 PHENOXYBENZAMINE HYDROCHLORIDE, PHENOXYBENZAMINE HYDROCHLORIDE
 PREDNISONE INTENSOL, PREDNISONE
 PREDNISONE, PREDNISONE
 PROPANTHELINE BROMIDE, PROPANTHELINE BROMIDE
 PROPRANOLOL HYDROCHLORIDE, PROPRANOLOL HYDROCHLORIDE
 PROTRIPTYLINE HYDROCHLORIDE, PROTRIPTYLINE HYDROCHLORIDE
 QUETIAPINE FUMARATE, QUETIAPINE FUMARATE
 RAMIPRIL, RAMIPRIL
 RISPERIDONE, RISPERIDONE
 RITONAVIR, RITONAVIR
 ROXICET, ACETAMINOPHEN
 RUFINAMIDE, RUFINAMIDE
 SODIUM OXYBATE, SODIUM OXYBATE
 TETRABENAZINE, TETRABENAZINE
 TINIDAZOLE, TINIDAZOLE
 TORSEMIDE, TORSEMIDE
 TRIAZOLAM, TRIAZOLAM
 VALACYCLOVIR HYDROCHLORIDE, VALACYCLOVIR HYDROCHLORIDE
 ZALEPLON, ZALEPLON
 ZIDOVUDINE, ZIDOVUDINE

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 DEXTROSE 5% IN PLASTIC CONTAINER, FLUCONAZOLE
 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
 CEFUROXIME SODIUM, CEFUROXIME SODIUM
 FLUCONAZOLE IN SODIUM CHLORIDE 0.9%, FLUCONAZOLE

* HIKMA FARMACEUTICA PORTUGAL SA

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** H ****

* HIKMA FARMACEUTICA PORTUGAL SA
 DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
 OXYTOCIN, OXYTOCIN
 TESTOSTERONE ENANTHATE, TESTOSTERONE ENANTHATE

* HIKMA FARMACEUTICA SA
 ZOLEDRONIC ACID, ZOLEDRONIC ACID

HIKMA INTL PHARMS

* HIKMA INTERNATIONAL PHARMACEUTICALS LLC
 BUTALBITAL, ASPIRIN AND CAFFEINE, ASPIRIN
 CAPTOPRIL, CAPTOPRIL
 CORTISONE ACETATE, CORTISONE ACETATE
 DIGOXIN, DIGOXIN
 DOPAMINE HYDROCHLORIDE, DOPAMINE HYDROCHLORIDE
 DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
 ERGOTAMINE TARTRATE AND CAFFEINE, CAFFEINE
 HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 HYDROCORTISONE, HYDROCORTISONE
 ISOSORBIDE DINITRATE, ISOSORBIDE DINITRATE
 LISINOPRIL AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 MITIGARE, COLCHICINE
 PRIMIDONE, PRIMIDONE

HIKMA PHARM CO LTD

* HIKMA PHARM CO LTD
 ARGATROBAN, ARGATROBAN

HIKMA PHARMS

* HIKMA PHARMACEUTICALS
 AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
 AMOXICILLIN AND CLAVULANATE POTASSIUM, AMOXICILLIN
 AMOXICILLIN, AMOXICILLIN
 CEFADROXIL, CEFADROXIL/CEFADROXIL HEMIHYDRATE
 DICYCLOMINE HYDROCHLORIDE, DICYCLOMINE HYDROCHLORIDE
 DIHYDROERGOTAMINE MESYLATE, DIHYDROERGOTAMINE MESYLATE
 ESCITALOPRAM OXALATE, ESCITALOPRAM OXALATE
 FLUDROCORTISONE ACETATE, FLUDROCORTISONE ACETATE
 GEMFIBROZIL, GEMFIBROZIL
 LETROZOLE, LETROZOLE
 PENICILLIN V POTASSIUM, PENICILLIN V POTASSIUM
 RIFAMPIN, RIFAMPIN
 VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE

* HIKMA PHARMACEUTICALS CO LTD
 PARICALCITOL, PARICALCITOL

* HIKMA PHARMACEUTICALS LLC
 ABIRATERONE ACETATE, ABIRATERONE ACETATE
 CHLOROQUINE PHOSPHATE, CHLOROQUINE PHOSPHATE
 DANTROLENE SODIUM, DANTROLENE SODIUM
 DOXERCALCIFEROL, DOXERCALCIFEROL
 FOLIC ACID, FOLIC ACID
 HYDROXYCHLOROQUINE SULFATE, HYDROXYCHLOROQUINE SULFATE
 ISOSORBIDE MONONITRATE, ISOSORBIDE MONONITRATE
 MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
 OLANZAPINE, OLANZAPINE
 PIROXICAM, PIROXICAM
 PREDNISONE, PREDNISONE
 ZALEPLON, ZALEPLON

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

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

**** H ****

* 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
IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE
LEVETIRACETAM, LEVETIRACETAM

* HISUN PHARMACEUTICAL HANGZHOU CO LTD
CAPREOMYCIN SULFATE, CAPREOMYCIN SULFATE
DACTINOMYCIN, DACTINOMYCIN
DAUNORUBICIN HYDROCHLORIDE, DAUNORUBICIN HYDROCHLORIDE
DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
PRAVASTATIN SODIUM, PRAVASTATIN SODIUM
TICAGRELOR, TICAGRELOR

HLTHCARE

* HEALTHCARE PHARMACEUTICALS LTD
ATENOLOL, ATENOLOL

HOFFMANN LA ROCHE

* HOFFMANN LA ROCHE INC
BONIVA, IBANDRONATE SODIUM
VALCYTE, VALGANCICLOVIR HYDROCHLORIDE
XELODA, CAPECITABINE
ZELBORAF, VEMURAFENIB

HOFFMANN-LA ROCHE

* HOFFMANN-LA ROCHE INC
ALECENSA, ALECTINIB HYDROCHLORIDE
INVIRASE, SAQUINAVIR MESYLATE

HOPE PHARMS

* HOPE PHARMACEUTICALS
NITHIODOTE, SODIUM NITRITE
SODIUM NITRITE, SODIUM NITRITE
SODIUM THIOSULFATE, SODIUM THIOSULFATE

HORIZON

* HORIZON MEDICINES LLC
DUEXIS, FAMOTIDINE
VIMOVO, ESOMEPRAZOLE MAGNESIUM

* HORIZON THERAPEUTICS IRELAND DAC
PENNSAID, DICLOFENAC SODIUM

HORIZON PHARMA USA

* HORIZON PHARMA USA INC
PROCYSBI, CYSTEAMINE BITARTRATE
RAYOS, PREDNISONE

HORIZON THERAP

* HORIZON THERAPEUTICS LLC
BUPHENYL, SODIUM PHENYLBUTYRATE
RAVICTI, GLYCEROL PHENYLBUTYRATE

HOSPIRA

* HOSPIRA INC
ACETYLCYSTEINE, ACETYLCYSTEINE
ALFENTANIL, ALFENTANIL HYDROCHLORIDE
AMIDATE, ETOMIDATE
AMINOCAPROIC ACID IN PLASTIC CONTAINER, AMINOCAPROIC ACID
AMINOPHYLLINE, AMINOPHYLLINE
AMMONIUM CHLORIDE IN PLASTIC CONTAINER, AMMONIUM CHLORIDE
ATROPINE SULFATE ANSYR PLASTIC SYRINGE, ATROPINE SULFATE
ATROPINE SULFATE LIFESHIELD ABBOJECT SYRINGE, ATROPINE SULFATE
AZITHROMYCIN, AZITHROMYCIN
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
BUMETANIDE, BUMETANIDE
BUPIVACAINE HYDROCHLORIDE AND EPINEPHRINE, BUPIVACAINE HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** H **

* HOSPIRA INC

BUPIVACAINE HYDROCHLORIDE, BUPIVACAINE HYDROCHLORIDE
 BUPRENORPHINE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE
 BUTORPHANOL TARTRATE PRESERVATIVE FREE, BUTORPHANOL TARTRATE
 CALCIUM CHLORIDE 10% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 CARBOCAINE, MEPIVACAINE HYDROCHLORIDE
 CARBOPLATIN, CARBOPLATIN
 CHLOROPROCAINE HYDROCHLORIDE, CHLOROPROCAINE HYDROCHLORIDE
 CHROMIC CHLORIDE IN PLASTIC CONTAINER, CHROMIC CHLORIDE
 CIPROFLOXACIN IN DEXTROSE 5% IN PLASTIC CONTAINER, CIPROFLOXACIN
 CIPROFLOXACIN, CIPROFLOXACIN
 CORLOPAM, FENOLDOPAM MESYLATE
 CUPRIC CHLORIDE IN PLASTIC CONTAINER, CUPRIC CHLORIDE
 CYTARABINE, CYTARABINE
 DACARBAZINE, DACARBAZINE
 DEFEROXAMINE MESYLATE, DEFEROXAMINE MESYLATE
 DEMEROL, MEPERIDINE HYDROCHLORIDE
 DEXTROSE 25%, DEXTROSE
 DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 50% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 50%, DEXTROSE
 DIAZEPAM, DIAZEPAM
 DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
 DIPHENHYDRAMINE HYDROCHLORIDE, DIPHENHYDRAMINE 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
 DROPERIDOL, DROPERIDOL
 ENALAPRILAT, ENALAPRILAT
 ERYTHROCIN, ERYTHROMYCIN LACTOBIONATE
 FENTANYL CITRATE PRESERVATIVE FREE, FENTANYL CITRATE
 FENTANYL CITRATE, FENTANYL CITRATE
 FLUCONAZOLE IN DEXTROSE 5% IN PLASTIC CONTAINER, FLUCONAZOLE
 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 20,000 UNITS IN DEXTROSE 5% IN PLASTIC CONTAINER, 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% 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
 LEVOPHED, NOREPINEPHRINE BITARTRATE
 LIDOCAINE HYDROCHLORIDE 5% AND DEXTROSE 7.5%, LIDOCAINE HYDROCHLORIDE
 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
 LORAZEPAM, LORAZEPAM
 M.V.I. ADULT (PHARMACY BULK PACKAGE), ASCORBIC ACID
 M.V.I. ADULT, ASCORBIC ACID
 M.V.I. PEDIATRIC, ASCORBIC ACID
 MAGNESIUM SULFATE IN DEXTROSE 5% IN PLASTIC CONTAINER, MAGNESIUM SULFATE
 MAGNESIUM SULFATE IN PLASTIC CONTAINER, MAGNESIUM SULFATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** H ****

* HOSPIRA INC
 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 IN DEXTROSE 5% IN PLASTIC CONTAINER, 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
 PAMIDRONATE DISODIUM, PAMIDRONATE DISODIUM
 PANCURONIUM BROMIDE, PANCURONIUM BROMIDE
 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
 ROCURONIUM BROMIDE, ROCURONIUM BROMIDE
 ROPIVACAINE HYDROCHLORIDE, ROPIVACAINE HYDROCHLORIDE
 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 IN PLASTIC CONTAINER, SODIUM CHLORIDE
 SODIUM LACTATE IN PLASTIC CONTAINER, SODIUM LACTATE
 SODIUM PHOSPHATES IN PLASTIC CONTAINER, SODIUM PHOSPHATE, DIBASIC, HEPTAHYDRATE
 STERILE WATER FOR INJECTION, STERILE WATER FOR INJECTION
 SUFENTANIL CITRATE, SUFENTANIL CITRATE
 TAZICEF, CEFTAZIDIME
 TOBRAMYCIN SULFATE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, TOBRAMYCIN SULFATE
 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

* HOSPIRA WORLDWIDE, INC
 DOBUTAMINE HYDROCHLORIDE, DOBUTAMINE HYDROCHLORIDE
 FLUDARABINE PHOSPHATE, FLUDARABINE PHOSPHATE
 GEMCITABINE HYDROCHLORIDE, GEMCITABINE HYDROCHLORIDE
 MITOXANTRONE HYDROCHLORIDE, MITOXANTRONE HYDROCHLORIDE
 NITROPRESS, SODIUM NITROPRUSSIDE
 TOBRAMYCIN SULFATE, TOBRAMYCIN SULFATE
 VINCRISTINE SULFATE PFS, VINCRISTINE SULFATE

HOSPIRA INC

* HOSPIRA INC
 ADENOSINE, ADENOSINE
 AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE
 ARGATROBAN, ARGATROBAN

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** H ******* HOSPIRA INC**

ATRACURIUM BESYLATE PRESERVATIVE FREE, ATRACURIUM BESYLATE
 ATRACURIUM BESYLATE, ATRACURIUM BESYLATE
 BIVALIRUDIN, BIVALIRUDIN
 BUSULFAN, BUSULFAN
 CISATRACURIUM BESYLATE, CISATRACURIUM BESYLATE
 DAPTOMYCIN, DAPTOMYCIN
 DOCETAXEL, DOCETAXEL
 DOXERCALCIFEROL, DOXERCALCIFEROL
 EPINEPHRINE (COPACKAGED), EPINEPHRINE
 GEMCITABINE HYDROCHLORIDE, GEMCITABINE HYDROCHLORIDE
 HEPARIN SODIUM, HEPARIN SODIUM
 HYDROMORPHONE HYDROCHLORIDE, HYDROMORPHONE HYDROCHLORIDE
 INDOMETHACIN SODIUM, INDOMETHACIN SODIUM
 LEVETIRACETAM, LEVETIRACETAM
 LEVOFLOXACIN IN DEXTROSE 5% IN PLASTIC CONTAINER, LEVOFLOXACIN
 LINEZOLID IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, LINEZOLID
 LINEZOLID, LINEZOLID
 MAGNESIUM SULFATE, MAGNESIUM SULFATE
 MAXIPIME, CEFEPIME HYDROCHLORIDE
 MILRINONE LACTATE, MILRINONE LACTATE
 MORPHINE SULFATE, MORPHINE SULFATE
 NIPENT, PENTOSTATIN
 PALONOSETRON HYDROCHLORIDE, PALONOSETRON HYDROCHLORIDE
 PARICALCITOL, PARICALCITOL
 SODIUM BICARBONATE, SODIUM BICARBONATE
 TACROLIMUS, TACROLIMUS
 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
 FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18
 SODIUM FLUORIDE F-18, SODIUM FLUORIDE F-18

HQ SPCLT PHARMA

* HQ SPECIALTY PHARMA CORP
 CALCIUM GLUCONATE IN SODIUM CHLORIDE, CALCIUM GLUCONATE
 CISPLATIN, CISPLATIN
 DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE
 ESMOLOL HYDROCHLORIDE DOUBLE STRENGTH IN PLASTIC CONTAINER, ESMOLOL HYDROCHLORIDE
 ESMOLOL HYDROCHLORIDE IN PLASTIC CONTAINER, ESMOLOL HYDROCHLORIDE
 IMIPENEM AND CILASTATIN, CILASTATIN SODIUM
 LINEZOLID, LINEZOLID
 MAGNESIUM SULFATE IN DEXTROSE 5% IN PLASTIC CONTAINER, MAGNESIUM SULFATE
 MAGNESIUM SULFATE IN PLASTIC CONTAINER, MAGNESIUM SULFATE
 MEROPENEM, MEROPENEM

HQ SPECIALITY PHARMA

* HQ SPECIALITY PHARMA LLC
 LEVETIRACETAM IN SODIUM CHLORIDE, LEVETIRACETAM

HRA PHARMA

* HRA PHARMA RARE DISEASES
 LYSODREN, MITOTANE
 METOPIRONE, METYRAPONE

HUMANWELL PURACAP

* HUMANWELL PURACAP PHARMACEUTICAL WUHAN CO LTD
 DUTASTERIDE, DUTASTERIDE
 IBUPROFEN, IBUPROFEN (OTC)

HUONS

* HUONS CO LTD
 BUPIVACAINE HYDROCHLORIDE, BUPIVACAINE HYDROCHLORIDE

ROCHE

* HOFFMANN LA ROCHE INC

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

**** H ****

* HOFFMANN LA ROCHE INC
 BONIVA, IBANDRONATE SODIUM
 FUZEON, ENFUVIRTIDE
 KLONOPIN, CLONAZEPAM
 TAMIFLU, OSELTAMIVIR PHOSPHATE
 VALIUM, DIAZEPAM

SHUANGCHENG

* HAINAN SHUANGCHENG PHARMACEUTICALS CO LTD
 BIVALIRUDIN, BIVALIRUDIN
 PREGABALIN, PREGABALIN

**** I ****

ICHNOS

* ICHNOS SCIENCES SA
 DEFERASIROX, DEFERASIROX

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
 MORPHINE SULFATE, MORPHINE SULFATE
 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
 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

IDENTI PHARMS INC

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** I ****

* IDENTI PHARMACEUTICALS INC
 FLUOCINOLONE ACETONIDE, FLUOCINOLONE ACETONIDE

IDT AUSTRALIA LTD

* IDT AUSTRALIA LTD
 TEMOZOLOMIDE, TEMOZOLOMIDE

IMPACT

* IMPACT BIOMEDICINES INC A WHOLLY OWNED SUB OF CELGENE CORP
 INREBIC, FEDRATINIB HYDROCHLORIDE

IMPAX

* IMPAX LABORATORIES LLC
 ADRENACLICK, EPINEPHRINE

IMPAX LABS

* IMPAX LABORATORIES INC
 ACARBOSE, ACARBOSE
 ANAGRELIDE HYDROCHLORIDE, ANAGRELIDE HYDROCHLORIDE
 BACLOFEN, BACLOFEN
 BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
 CARBIDOPA AND LEVODOPA, CARBIDOPA
 CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
 COLESTIPOL HYDROCHLORIDE, COLESTIPOL HYDROCHLORIDE
 DANTROLENE SODIUM, DANTROLENE SODIUM
 DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE
 DIGOXIN, DIGOXIN
 DIPYRIDAMOLE, DIPYRIDAMOLE
 DIVALPROEX SODIUM, DIVALPROEX SODIUM
 FENOFIBRATE (MICRONIZED), FENOFIBRATE
 FENOFIBRATE, FENOFIBRATE
 FLUDROCORTISONE ACETATE, FLUDROCORTISONE ACETATE
 METHYLTESTOSTERONE, METHYLTESTOSTERONE
 MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE
 NADOLOL AND BENDROFLUMETHIAZIDE, BENDROFLUMETHIAZIDE
 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
 ALBENZA, ALBENDAZOLE
 ALENDRONATE SODIUM, ALENDRONATE SODIUM
 BUDESONIDE, BUDESONIDE
 BUSPIRONE HYDROCHLORIDE, BUSPIRONE HYDROCHLORIDE
 CARVEDILOL PHOSPHATE, CARVEDILOL PHOSPHATE
 COLESEVELAM HYDROCHLORIDE, COLESEVELAM HYDROCHLORIDE
 DESMOPRESSIN ACETATE, DESMOPRESSIN ACETATE
 DEXEDRINE, DEXTROAMPHETAMINE SULFATE
 DEXMETHYLPHENIDATE HYDROCHLORIDE, DEXMETHYLPHENIDATE HYDROCHLORIDE
 DOXYCYCLINE, DOXYCYCLINE
 EMVERM, MEBENDAZOLE
 EPIRUBICIN HYDROCHLORIDE, EPIRUBICIN HYDROCHLORIDE
 FENOFIBRIC ACID, CHOLINE FENOFIBRATE
 GLYBURIDE AND METFORMIN HYDROCHLORIDE, GLYBURIDE
 GLYBURIDE, GLYBURIDE
 HYDROCORTISONE, HYDROCORTISONE
 HYDROXYZINE PAMOATE, HYDROXYZINE PAMOATE
 LAMOTRIGINE, LAMOTRIGINE
 LEVALBUTEROL HYDROCHLORIDE, LEVALBUTEROL HYDROCHLORIDE
 METHENAMINE HIPPURATE, METHENAMINE HIPPURATE
 METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
 METHYLTESTOSTERONE, METHYLTESTOSTERONE
 METOCLOPRAMIDE HYDROCHLORIDE, METOCLOPRAMIDE HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

**** I ****

* IMPAX LABORATORIES INC
 MIRTAZAPINE, MIRTAZAPINE
 MORPHINE SULFATE, MORPHINE SULFATE
 NABUMETONE, NABUMETONE
 NITROFURANTOIN, NITROFURANTOIN, MACROCRYSTALLINE
 PROPRANOLOL HYDROCHLORIDE, PROPRANOLOL HYDROCHLORIDE
 PYRIDOSTIGMINE BROMIDE, PYRIDOSTIGMINE BROMIDE
 RYTARY, CARBIDOPA
 SEVELAMER CARBONATE, SEVELAMER CARBONATE
 URSODIOL, URSODIOL

IMPAX PHARMS

* IMPAX PHARMACEUTICALS
 GEMFIBROZIL, GEMFIBROZIL
 MIDODRINE HYDROCHLORIDE, MIDODRINE HYDROCHLORIDE
 ORPHENADRINE CITRATE, ORPHENADRINE CITRATE
 OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE

INCYTE CORP

* INCYTE CORP
 JAKAFI, RUXOLITINIB PHOSPHATE

INDICUS PHARMA

* INDICUS PHARMA LLC
 DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
 LETROZOLE, LETROZOLE
 MECLIZINE HYDROCHLORIDE, MECLIZINE HYDROCHLORIDE
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE

INDIVIOR INC

* INDIVIOR INC
 BUPRENEX, BUPRENORPHINE HYDROCHLORIDE
 PERSERIS KIT, RISPERIDONE
 SUBLOCADE, BUPRENORPHINE
 SUBOXONE, BUPRENORPHINE HYDROCHLORIDE

INDOCO REMEDIES

* INDOCO REMEDIES LTD
 ALLOPURINOL, ALLOPURINOL
 BRIMONIDINE TARTRATE, BRIMONIDINE TARTRATE
 FEBUXOSTAT, FEBUXOSTAT
 GLIMEPIRIDE, GLIMEPIRIDE
 RASAGILINE MESYLATE, RASAGILINE MESYLATE

INFORLIFE

* INFORLIFE SA
 CIPROFLOXACIN IN DEXTROSE 5% IN PLASTIC CONTAINER, CIPROFLOXACIN
 FLUCONAZOLE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, FLUCONAZOLE
 LEVOFLOXACIN IN DEXTROSE 5% IN PLASTIC CONTAINER, LEVOFLOXACIN
 METRONIDAZOLE IN PLASTIC CONTAINER, METRONIDAZOLE
 ROPIVACAINE HYDROCHLORIDE, ROPIVACAINE HYDROCHLORIDE
 ZOLEDRONIC ACID, ZOLEDRONIC ACID

INGENUS PHARMS LLC

* INGENUS PHARMACEUTICALS LLC
 ARSENIC TRIOXIDE, ARSENIC TRIOXIDE
 CABERGOLINE, CABERGOLINE
 CARBOPLATIN, CARBOPLATIN
 CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
 CLOFARABINE, CLOFARABINE
 DECITABINE, DECITABINE
 DOCETAXEL, DOCETAXEL
 FLUOROURACIL, FLUOROURACIL
 IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE
 LEVOLEUCOVORIN CALCIUM, LEVOLEUCOVORIN CALCIUM
 MELPHALAN HYDROCHLORIDE, MELPHALAN HYDROCHLORIDE
 OXALIPLATIN, OXALIPLATIN
 PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
 ZOLEDRONIC ACID, ZOLEDRONIC ACID

INGENUS PHARMS NJ

* INGENUS PHARMACEUTICALS NJ LLC
 CARISOPRODOL, ASPIRIN AND CODEINE PHOSPHATE, ASPIRIN

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** I ******INNOGENIX**

- * INNOGENIX LLC
 - HALOPERIDOL, HALOPERIDOL
 - LABETALOL HYDROCHLORIDE, LABETALOL HYDROCHLORIDE
 - METRONIDAZOLE, METRONIDAZOLE
 - PILOCARPINE HYDROCHLORIDE, PILOCARPINE HYDROCHLORIDE
 - PROPRANOLOL HYDROCHLORIDE, PROPRANOLOL HYDROCHLORIDE

INSMED INC

- * INSMED INC
 - ARIKAYCE KIT, AMIKACIN SULFATE

INST BIOCHEM

- * INSTITUT BIOCHEMIQUE SA
 - FLECTOR, DICLOFENAC EPOLAMINE

INSTITUT BIOCHIMIQUE

- * INSTITUT BIOCHIMIQUE SA (IBSA)
 - TIROSINT, LEVOTHYROXINE SODIUM
 - TIROSINT-SOL, LEVOTHYROXINE SODIUM

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
 - OALIVA, OBETICHOLIC ACID

INTERGEL PHARM

- * INTERGEL PHARMACEUTICAL INC
 - NIFEDIPINE, NIFEDIPINE

INTERGEL PHARMS INC

- * INTERGEL PHARMACEUTICALS INC
 - DUTASTERIDE, DUTASTERIDE

INTERPHARMA PRAHA AS

- * INTERPHARMA PRAHA AS
 - ORALTAG, IOHEXOL

INTERSECT ENT INC

- * INTERSECT ENT INC
 - SINUVA, MOMETASONE FUROATE

INTL ISOTOPES

- * INTERNATIONAL ISOTOPES INC
 - SODIUM IODIDE I 131, SODIUM IODIDE I-131

INTL MEDICATED

- * INTERNATIONAL MEDICATED SYSTEMS LTD
 - MILRINONE LACTATE, MILRINONE LACTATE

INTL MEDICATION

- * INTERNATIONAL MEDICATION SYSTEM
 - LARYNG-O-JET KIT, LIDOCAINE HYDROCHLORIDE
 - LIDOCAINE HYDROCHLORIDE, LIDOCAINE HYDROCHLORIDE
 - MANNITOL 25%, MANNITOL
 - NALOXONE HYDROCHLORIDE, NALOXONE HYDROCHLORIDE
 - PHYTONADIONE, PHYTONADIONE
 - PROCAINAMIDE HYDROCHLORIDE, PROCAINAMIDE HYDROCHLORIDE
- * INTERNATIONAL MEDICATION SYSTEMS LTD
 - DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE

INTL MEDICATION SYS

- * INTERNATIONAL MEDICATION SYSTEMS LTD
 - CALCIUM CHLORIDE 10%, CALCIUM CHLORIDE
 - LORAZEPAM, LORAZEPAM
 - SODIUM BICARBONATE, SODIUM BICARBONATE

INTRA-CELLULAR

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

**** I ****

* INTRA-CELLULAR THERAPIES INC
CAPLYTA, LUMATEPERONE TOSYLATE

INVAGEN PHARMS

* INVAGEN PHARMACEUTICALS INC
ALFUZOSIN HYDROCHLORIDE, ALFUZOSIN HYDROCHLORIDE
AMANTADINE HYDROCHLORIDE, AMANTADINE HYDROCHLORIDE
AMLODIPINE BESYLATE AND VALSARTAN, AMLODIPINE BESYLATE
AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM
BENZTROPINE MESYLATE, BENZTROPINE MESYLATE
BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
CALCIUM ACETATE, CALCIUM ACETATE
CARBINOXAMINE MALEATE, CARBINOXAMINE MALEATE
CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE
CYCLOBENZAPRINE HYDROCHLORIDE, CYCLOBENZAPRINE HYDROCHLORIDE
ESCITALOPRAM OXALATE, ESCITALOPRAM OXALATE
FENOFIBRATE (MICRONIZED), FENOFIBRATE
FOSINOPRIL SODIUM AND HYDROCHLOROTHIAZIDE, FOSINOPRIL SODIUM
FOSINOPRIL SODIUM, FOSINOPRIL SODIUM
GABAPENTIN, GABAPENTIN
GEMFIBROZIL, GEMFIBROZIL
GLIMEPIRIDE, GLIMEPIRIDE
HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE
LEVETIRACETAM, LEVETIRACETAM
LISINOPRIL AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
LISINOPRIL, LISINOPRIL
MEPROBAMATE, MEPROBAMATE
NABUMETONE, NABUMETONE
NADOLOL, NADOLOL
NAPROXEN, NAPROXEN
OLANZAPINE, OLANZAPINE
ORPHENADRINE CITRATE, ORPHENADRINE CITRATE
PHENTERMINE HYDROCHLORIDE, PHENTERMINE HYDROCHLORIDE
PREGABALIN, PREGABALIN
QUINAPRIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
QUINAPRIL HYDROCHLORIDE, QUINAPRIL HYDROCHLORIDE
RALOXIFENE HYDROCHLORIDE, RALOXIFENE HYDROCHLORIDE
RIZATRIPTAN BENZOATE, RIZATRIPTAN BENZOATE
SERTRALINE HYDROCHLORIDE, SERTRALINE HYDROCHLORIDE
SEVELAMER CARBONATE, SEVELAMER CARBONATE
TERBINAFINE HYDROCHLORIDE, TERBINAFINE HYDROCHLORIDE
TOPIRAMATE, TOPIRAMATE
TROSPIMUM CHLORIDE, TROSPIMUM CHLORIDE
VIGABATRIN, VIGABATRIN
WARFARIN SODIUM, WARFARIN SODIUM
ZOLMITRIPTAN, ZOLMITRIPTAN
ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE
ZONISAMIDE, ZONISAMIDE

INVATECH

* INVATECH PHARMA SOLUTIONS LLC
AMANTADINE HYDROCHLORIDE, AMANTADINE HYDROCHLORIDE
DIVALPROEX SODIUM, DIVALPROEX SODIUM
MECLIZINE HYDROCHLORIDE, MECLIZINE HYDROCHLORIDE
PYRIDOSTIGMINE BROMIDE, PYRIDOSTIGMINE BROMIDE
TOLTERODINE TARTRATE, TOLTERODINE TARTRATE

INVENTIA

* INVENTIA HEALTHCARE LTD
DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE
LANSOPRAZOLE, LANSOPRAZOLE
METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
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

IPCA LABS LTD

* IPCA LABORATORIES LTD
 ALLOPURINOL, ALLOPURINOL
 ATENOLOL, ATENOLOL
 CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 CETIRIZINE HYDROCHLORIDE HIVES, CETIRIZINE HYDROCHLORIDE (OTC)
 CHLOROQUINE PHOSPHATE, CHLOROQUINE PHOSPHATE
 FUROSEMIDE, FUROSEMIDE
 HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 HYDROXYCHLOROQUINE SULFATE, HYDROXYCHLOROQUINE SULFATE
 LOSARTAN POTASSIUM AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 LOSARTAN POTASSIUM, LOSARTAN POTASSIUM
 METOCLOPRAMIDE HYDROCHLORIDE, METOCLOPRAMIDE HYDROCHLORIDE
 METOPROLOL TARTRATE, METOPROLOL TARTRATE
 ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
 PROPRANOLOL HYDROCHLORIDE, PROPRANOLOL HYDROCHLORIDE
 TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE
 WARFARIN SODIUM, WARFARIN SODIUM

IPR

* IPR PHARMACEUTICALS INC
 CRESTOR, ROSUVASTATIN CALCIUM
 ZOMIG, ZOLMITRIPTAN

IPSEN INC

* IPSEN BIOPHARMACEUTICALS INC
 INCRELEX, MECASERMIN RECOMBINANT
 ONIVYDE, IRINOTECAN HYDROCHLORIDE

IPSEN PHARMA

* IPSEN PHARMA BIOTECH SAS
 SOMATULINE DEPOT, LANREOTIDE ACETATE

IRONSHORE PHARMS

* IRONSHORE PHARMACEUTICALS AND DEVELOPMENT INC
 JORNAY PM, METHYLPHENIDATE HYDROCHLORIDE

ISO TEX

* ISO TEX DIAGNOSTICS INC
 JEANATOPE, ALBUMIN IODINATED I-125 SERUM
 MEGATOPE, ALBUMIN IODINATED I-131 SERUM

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 SPA

* ITALFARMACO SPA
 TIGLUTIK KIT, RILUZOLE

IVAX PHARMS

* IVAX PHARMACEUTICALS INC
 VALSARTAN, VALSARTAN

IVAX PHARMS INC

* IVAX PHARMACEUTICALS INC
 OLANZAPINE, OLANZAPINE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** I ******IVAX SUB TEVA PHARMS**

* IVAX PHARMACEUTICALS INC SUB TEVA PHARMACEUTICALS USA
 ANAGRELIDE HYDROCHLORIDE, ANAGRELIDE HYDROCHLORIDE
 BACLOFEN, BACLOFEN
 CABERGOLINE, CABERGOLINE
 CETIRIZINE HYDROCHLORIDE AND PSEUDOEPHEDRINE HYDROCHLORIDE, CETIRIZINE HYDROCHLORIDE
 CIMETIDINE, CIMETIDINE (OTC)
 CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE
 CLOZAPINE, CLOZAPINE
 CYCLOSPORINE, CYCLOSPORINE
 DIAZEPAM, DIAZEPAM
 ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM
 FAMOTIDINE, FAMOTIDINE
 FLUCONAZOLE, FLUCONAZOLE
 FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 GABAPENTIN, GABAPENTIN
 HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 OXAPROZIN, OXAPROZIN
 TOLTERODINE TARTRATE, TOLTERODINE TARTRATE
 VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE

**** J ******J AND J CONSUMER INC**

* JOHNSON AND JOHNSON CONSUMER INC MCNEIL CONSUMER HEALTHCARE DIVISION
 CHILDREN'S MOTRIN COLD, IBUPROFEN (OTC)
 CHILDREN'S MOTRIN, IBUPROFEN (OTC)
 CHILDREN'S ZYRTEC ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 CHILDREN'S ZYRTEC HIVES RELIEF, CETIRIZINE HYDROCHLORIDE (OTC)
 IMODIUM A-D, LOPERAMIDE HYDROCHLORIDE (OTC)
 IMODIUM MULTI-SYMPTOM RELIEF, LOPERAMIDE HYDROCHLORIDE (OTC)
 JUNIOR STRENGTH MOTRIN, IBUPROFEN (OTC)
 MOTRIN IB, IBUPROFEN (OTC)
 PEPCID AC, FAMOTIDINE (OTC)
 PEPCID COMPLETE, CALCIUM CARBONATE (OTC)
 SINE-AID IB, IBUPROFEN (OTC)
 SUDAFED 24 HOUR, PSEUDOEPHEDRINE HYDROCHLORIDE (OTC)
 TYLENOL, ACETAMINOPHEN (OTC)
 ZYRTEC ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 ZYRTEC-D 12 HOUR, CETIRIZINE HYDROCHLORIDE (OTC)

JACOBUS

* JACOBUS PHARMACEUTICAL CO
 DAPSONE, DAPSONE
 PASER, AMINOSALICYLIC ACID

JACOBUS PHARM CO INC

* JACOBUS PHARMACEUTICAL CO INC
 RUZURGI, AMIFAMPRIDINE

JANSSEN BIOTECH

* JANSSEN BIOTECH INC
 BALVERSA, ERDAFITINIB
 ERLEADA, APALUTAMIDE
 ZYTIGA, ABIRATERONE ACETATE

JANSSEN PHARMS

* JANSSEN PHARMACEUTICALS INC
 CONCERTA, METHYLPHENIDATE HYDROCHLORIDE
 DITROPAN XL, OXYBUTYNIN CHLORIDE
 DURAGESIC-100, FENTANYL
 DURAGESIC-12, FENTANYL
 DURAGESIC-25, FENTANYL
 DURAGESIC-37, FENTANYL
 DURAGESIC-50, FENTANYL
 DURAGESIC-75, FENTANYL
 ELMIRON, PENTOSAN POLYSULFATE SODIUM
 HALDOL, HALOPERIDOL DECANOATE
 HALDOL, HALOPERIDOL LACTATE
 INVEGA SUSTENNA, PALIPERIDONE PALMITATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** J ****

* JANSSEN PHARMACEUTICALS INC
 INVEGA TRINZA, PALIPERIDONE PALMITATE
 INVEGA, PALIPERIDONE
 INVOKAMET XR, CANAGLIFLOZIN
 INVOKAMET, CANAGLIFLOZIN
 INVOKANA, CANAGLIFLOZIN
 NIZORAL, KETOCONAZOLE
 ORTHO TRI-CYCLEN LO, ETHINYL ESTRADIOL
 ORTHO TRI-CYCLEN, ETHINYL ESTRADIOL
 ORTHO-NOVUM 7/7/7-28, ETHINYL ESTRADIOL
 RAZADYNE ER, GALANTAMINE HYDROBROMIDE
 RAZADYNE, GALANTAMINE HYDROBROMIDE
 RISPERDAL CONSTA, RISPERIDONE
 RISPERDAL, RISPERIDONE
 SPORANOX, ITRACONAZOLE
 SPRAVATO, ESKETAMINE HYDROCHLORIDE
 TOPAMAX, TOPIRAMATE
 TYLENOL W/ CODEINE NO. 3, ACETAMINOPHEN
 TYLENOL W/ CODEINE NO. 4, ACETAMINOPHEN
 ULTRACET, ACETAMINOPHEN
 ULTRAM, TRAMADOL HYDROCHLORIDE
 XARELTO, RIVAROXABAN

JANSSEN PRODS

* 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
 SUNOSI, SOLRIAMFETOL HYDROCHLORIDE

JAZZ PHARMS

* JAZZ PHARMACEUTICALS INC
 XYREM, SODIUM OXYBATE

JAZZ PHARMS INC

* JAZZ PHARMACEUTICALS INC
 DEFITELIO, DEFIBROTIDE SODIUM

JDP

* JDP THERAPEUTICS LLC
 QUZYTIR, CETIRIZINE HYDROCHLORIDE

JIANGSU HANSOH PHARM

* JIANGSU HANSOH PHARMACEUTICAL GROUP CO LTD
 GEMCITABINE HYDROCHLORIDE, GEMCITABINE HYDROCHLORIDE
 ICATIBANT ACETATE, ICATIBANT ACETATE
 OLANZAPINE, OLANZAPINE
 VINORELBINE TARTRATE, VINORELBINE TARTRATE

JIANGSU HENGRUI MED

* JIANGSU HENGRUI MEDICINE CO LTD
 CASPOFUNGIN ACETATE, CASPOFUNGIN ACETATE
 CISATRACURIUM BESYLATE PRESERVATIVE FREE, CISATRACURIUM BESYLATE
 CISATRACURIUM BESYLATE, CISATRACURIUM BESYLATE
 CYCLOPHOSPHAMIDE, CYCLOPHOSPHAMIDE
 DAPTOMYCIN, DAPTOMYCIN
 DEXMEDETOMIDINE, DEXMEDETOMIDINE HYDROCHLORIDE
 DOCETAXEL, DOCETAXEL
 FONDAPARINUX SODIUM, FONDAPARINUX SODIUM
 GABAPENTIN, GABAPENTIN
 IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** J ****

- * JIANGSU HENGRUI MEDICINE CO LTD
LETROZOLE, LETROZOLE
OXALIPLATIN, OXALIPLATIN
THIOTEPA, THIOTEPA

JIANGXI BOYA SEEHOT

- * JIANGXI BOYA SEEHOT PHARMACEUTICAL CO LTD
SOLIFENACIN SUCCINATE, SOLIFENACIN SUCCINATE

JOHNS HOPKINS UNIV

- * JOHNS HOPKINS UNIV
AMMONIA N 13, AMMONIA N-13

JOHNSON AND JOHNSON

- * JOHNSON AND JOHNSON CONSUMER INC
VISINE L.R., OXYMETAZOLINE HYDROCHLORIDE (OTC)
VISINE, NAPHAZOLINE HYDROCHLORIDE (OTC)
- * JOHNSON AND JOHNSON GROUP CONSUMER COMPANIES
MEN'S ROGAINE, MINOXIDIL (OTC)
ROGAINE (FOR MEN), MINOXIDIL (OTC)
ROGAINE (FOR WOMEN), MINOXIDIL (OTC)
ROGAINE EXTRA STRENGTH (FOR MEN), MINOXIDIL (OTC)
WOMEN'S ROGAINE, MINOXIDIL (OTC)

JOURNEY

- * JOURNEY MEDICAL CORP
EXELDERM, SULCONAZOLE NITRATE
XIMINO, MINOCYCLINE HYDROCHLORIDE

JUBILANT CADISTA

- * JUBILANT CADISTA PHARMACEUTICALS INC
ALENDRONATE SODIUM, ALENDRONATE SODIUM
CLOMIPRAMINE HYDROCHLORIDE, CLOMIPRAMINE HYDROCHLORIDE
CYCLOBENZAPRINE HYDROCHLORIDE, CYCLOBENZAPRINE HYDROCHLORIDE
DROSPIRENONE AND ETHINYL ESTRADIOL, DROSPIRENONE
HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
LAMOTRIGINE, LAMOTRIGINE
MECLIZINE HYDROCHLORIDE, MECLIZINE HYDROCHLORIDE
METHYLPREDNISOLONE, METHYLPREDNISOLONE
PREDNISONE, PREDNISONE
PROCOMP, PROCHLORPERAZINE MALEATE
SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, SODIUM CHLORIDE
TERAZOSIN HYDROCHLORIDE, TERAZOSIN HYDROCHLORIDE

JUBILANT DRAXIMAGE

- * JUBILANT DRAXIMAGE INC
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
- * 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
AMANTADINE HYDROCHLORIDE, AMANTADINE HYDROCHLORIDE
AMLODIPINE AND OLMESARTAN MEDOXOMIL, AMLODIPINE BESYLATE
CELECOXIB, CELECOXIB
CHILDREN'S CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
CHILDREN'S CETIRIZINE HYDROCHLORIDE HIVES RELIEF, CETIRIZINE HYDROCHLORIDE (OTC)
CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE
CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
DARIFENACIN HYDROBROMIDE, DARIFENACIN HYDROBROMIDE
DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
ESCITALOPRAM OXALATE, ESCITALOPRAM OXALATE
FELODIPINE, FELODIPINE
INDOMETHACIN, INDOMETHACIN

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** J ****

* JUBILANT GENERICS LTD
 IRBESARTAN, IRBESARTAN
 ITRACONAZOLE, ITRACONAZOLE
 LAMOTRIGINE, LAMOTRIGINE
 LEVETIRACETAM, LEVETIRACETAM
 LEVOFLOXACIN, LEVOFLOXACIN
 MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
 NIACIN, NIACIN
 OLANZAPINE, OLANZAPINE
 PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
 QUETIAPINE FUMARATE, QUETIAPINE FUMARATE
 RISPERIDONE, RISPERIDONE
 RIZATRIPTAN BENZOATE, RIZATRIPTAN BENZOATE
 ROSUVASTATIN CALCIUM, ROSUVASTATIN CALCIUM
 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
 METHOCARBAMOL AND ASPIRIN, ASPIRIN
 UNITHROID, LEVOTHYROXINE SODIUM **

**** K ******GRIFFEN**

* KW GRIFFEN CO
 BIOSCRUB, CHLORHEXIDINE GLUCONATE (OTC)

KADMON PHARMS LLC

* KADMON PHARMACEUTICALS LLC
 CLOVIQUE, TRIENTINE HYDROCHLORIDE
 TRIENTINE HYDROCHLORIDE, TRIENTINE HYDROCHLORIDE

KAI PHARMS INC

* KAI PHARMACEUTICALS INC A WHOLLY OWNED SUBSIDIARY OF AMGEN INC
 PARSABIV, ETELCALCETIDE

KALA PHARMS INC

* KALA PHARMACEUTICALS INC
 INVELTYS, LOTEPIREDNOL ETABONATE

KALEO INC

* KALEO INC
 AUVI-Q, EPINEPHRINE
 EVZIO (AUTOINJECTOR), NALOXONE HYDROCHLORIDE

KARYOPHARM THERAPS

* KARYOPHARM THERAPEUTICS INC
 XPOVIO, SELINEXOR

KENTON

* KENTON CHEMICALS AND PHARMACEUTICALS CORP
 ACYCLOVIR, ACYCLOVIR
 ANASTROZOLE, ANASTROZOLE
 BICALUTAMIDE, BICALUTAMIDE
 GLYCOPYRROLATE, GLYCOPYRROLATE

KERYX BIOPHARMS

* KERYX BIOPHARMACEUTICALS INC
 AURYXIA, FERRIC CITRATE

KETTERING MEDCTR

* KETTERING MEDCTR
 FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18

KING PHARMS

* KING PHARMACEUTICALS INC
 SYNERCID, DALFOPRISTIN
 * KING PHARMACEUTICALS RESEARCH AND DEVELOPMENT LLC
 CYTOMEL, LIOTHYRONINE SODIUM

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** K ****

- * KING PHARMACEUTICALS RESEARCH AND DEVELOPMENT LLC
LEVOXYL, LEVOTHYROXINE SODIUM **
- * KING PHARMACEUTICALS RESEARCH AND DEVELOPMENT LLC A SUB OF PFIZER INC
SKELAXIN, METAXALONE

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
CORZIDE, BENDROFLUMETHIAZIDE
PENICILLIN G PROCAINE, PENICILLIN G PROCAINE
SILVADENE, SILVER SULFADIAZINE
TAPAZOLE, METHIMAZOLE
TIGAN, TRIMETHOBENZAMIDE HYDROCHLORIDE

KNIGHT THERAPS

- * KNIGHT THERAPEUTICS USA INC
IMPAVIDO, MILTEFOSINE

KOWA CO

- * KOWA CO LTD
LIVALO, PITAVASTATIN CALCIUM

KRAMER

- * KRAMER LABORATORIES INC
NIZORAL A-D, 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
APADAZ, ACETAMINOPHEN
PHENTERMINE HYDROCHLORIDE, PHENTERMINE HYDROCHLORIDE

KYOWA KIRIN

- * KYOWA KIRIN INC
FARESTON, TOREMIFENE CITRATE
NOURIANZ, ISTRADefylline
SANCUSO, GRANISETRON

KYTHERA BIOPHARMS

- * KYTHERA BIOPHARMACEUTICALS INC
KYBELLA, DEOXYCHOLIC ACID

**** L ******L PERRIGO CO**

- * L PERRIGO CO
CIMETIDINE, CIMETIDINE (OTC)
CLEMASTINE FUMARATE, CLEMASTINE FUMARATE (OTC)
FEXOFENADINE HYDROCHLORIDE, FEXOFENADINE HYDROCHLORIDE (OTC)
IBUPROFEN, IBUPROFEN
IBUPROFEN, IBUPROFEN (OTC)
JUNIOR STRENGTH IBUPROFEN, IBUPROFEN (OTC)
LEVOCETIRIZINE DIHYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE
LEVONORGESTREL, LEVONORGESTREL
LEVONORGESTREL, LEVONORGESTREL (OTC)

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** L ******* L PERRIGO CO**

LOPERAMIDE HYDROCHLORIDE, LOPERAMIDE HYDROCHLORIDE (OTC)
 MINOXIDIL (FOR MEN), MINOXIDIL (OTC)
 MINOXIDIL (FOR WOMEN), MINOXIDIL (OTC)
 MONTELUKAST SODIUM, MONTELUKAST SODIUM
 NICOTINE POLACRILEX, NICOTINE POLACRILEX (OTC)
 PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
 PSEUDOEPHEDRINE HYDROCHLORIDE, PSEUDOEPHEDRINE HYDROCHLORIDE (OTC)

LA JOLLA PHARMA

* 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)

LABORATORIOS GRIFOLS

* LABORATORIOS GRIFOLS SA
 SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, SODIUM CHLORIDE

LABORATORIOS SALVAT

* LABORATORIOS SALVAT SA
 OTOVEL, CIPROFLOXACIN HYDROCHLORIDE

LANDELA PHARM

* LANDELA PHARMACEUTICAL
 FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 IPRATROPIUM BROMIDE, IPRATROPIUM BROMIDE

LANNETT

* LANNETT CO INC
 ACETAZOLAMIDE, ACETAZOLAMIDE
 DICYCLOMINE HYDROCHLORIDE, DICYCLOMINE HYDROCHLORIDE
 DIPHENOXYLATE HYDROCHLORIDE AND ATROPINE SULFATE, ATROPINE SULFATE
 LANORINAL, ASPIRIN
 PHENTERMINE HYDROCHLORIDE, PHENTERMINE HYDROCHLORIDE
 PRIMIDONE, PRIMIDONE
 PROBALAN, PROBENECID

LANNETT CO INC

* LANNETT CO INC
 ALBUTEROL SULFATE, ALBUTEROL SULFATE
 ARIPIPRAZOLE, ARIPIPRAZOLE
 ASPIRIN AND DIPYRIDAMOLE, ASPIRIN
 ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM
 BACLOFEN, BACLOFEN
 BETHANECHOL CHLORIDE, BETHANECHOL CHLORIDE
 BUPRENORPHINE HYDROCHLORIDE AND NALOXONE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE
 BUTALBITAL, ACETAMINOPHEN AND CAFFEINE, ACETAMINOPHEN
 CETIRIZINE HYDROCHLORIDE, CETIRIZINE HYDROCHLORIDE
 CHILDREN'S CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 CHILDREN'S CETIRIZINE HYDROCHLORIDE HIVES RELIEF, CETIRIZINE HYDROCHLORIDE (OTC)
 CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE
 CLINDAMYCIN HYDROCHLORIDE, CLINDAMYCIN HYDROCHLORIDE
 CLOBAZAM, CLOBAZAM
 CODEINE SULFATE, CODEINE SULFATE
 CYPROHEPTADINE HYDROCHLORIDE, CYPROHEPTADINE HYDROCHLORIDE
 DANAZOL, DANAZOL
 DEXAMETHASONE, DEXAMETHASONE
 DIAZEPAM, DIAZEPAM
 DIETHYLPROPION HYDROCHLORIDE, DIETHYLPROPION HYDROCHLORIDE
 DOXEPIN HYDROCHLORIDE, DOXEPIN HYDROCHLORIDE
 DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
 DOXYCYCLINE, DOXYCYCLINE
 DRONABINOL, DRONABINOL
 ESCITALOPRAM OXALATE, ESCITALOPRAM OXALATE
 ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM
 FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** L ****

* LANNETT CO INC
 FLUPHENAZINE HYDROCHLORIDE, FLUPHENAZINE HYDROCHLORIDE
 GLYCOLAX, POLYETHYLENE GLYCOL 3350 (OTC)
 HALOPERIDOL, HALOPERIDOL LACTATE
 HYDROCORTISONE, HYDROCORTISONE
 HYDROMORPHONE HYDROCHLORIDE, HYDROMORPHONE HYDROCHLORIDE
 HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE
 ISOSORBIDE MONONITRATE, ISOSORBIDE MONONITRATE
 LACTULOSE, LACTULOSE
 LAMIVUDINE, LAMIVUDINE
 LANSOPRAZOLE, LANSOPRAZOLE
 LANSOPRAZOLE, LANSOPRAZOLE (OTC)
 LEVETIRACETAM, LEVETIRACETAM
 LEVOCETIRIZINE DIHYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE
 LEVOFLOXACIN, LEVOFLOXACIN
 LIDOCAINE HYDROCHLORIDE VISCOUS, LIDOCAINE HYDROCHLORIDE
 LIDOCAINE HYDROCHLORIDE, LIDOCAINE HYDROCHLORIDE
 LOPINAVIR AND RITONAVIR, LOPINAVIR
 LORATADINE, LORATADINE (OTC)
 LORAZEPAM, LORAZEPAM
 LOXAPINE SUCCINATE, LOXAPINE SUCCINATE
 MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
 METADATE CD, METHYLPHENIDATE HYDROCHLORIDE
 METAPROTERENOL SULFATE, METAPROTERENOL SULFATE
 METAXALONE, METAXALONE
 METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
 MONOKET, ISOSORBIDE MONONITRATE
 MONTELUKAST SODIUM, MONTELUKAST SODIUM
 NIACIN, NIACIN
 NYSTATIN, NYSTATIN
 OMEPRAZOLE, OMEPRAZOLE
 ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
 OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE
 OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN
 OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
 PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
 PAROXETINE HYDROCHLORIDE, PAROXETINE HYDROCHLORIDE
 PHENTERMINE HYDROCHLORIDE, PHENTERMINE HYDROCHLORIDE
 PILOCARPINE HYDROCHLORIDE, PILOCARPINE HYDROCHLORIDE
 POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
 PREDNISOLONE, PREDNISOLONE
 RABEPRAZOLE SODIUM, RABEPRAZOLE SODIUM
 RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
 RIFAMPIN, RIFAMPIN
 RISPERIDONE, RISPERIDONE
 SULFAMETHOXAZOLE AND TRIMETHOPRIM, SULFAMETHOXAZOLE
 SUMATRIPTAN, SUMATRIPTAN
 TERBUTALINE SULFATE, TERBUTALINE SULFATE
 THEOPHYLLINE, THEOPHYLLINE
 TRIAMTERENE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 URSODIOL, URSODIOL
 VALPROIC ACID, VALPROIC ACID
 ZAROXOLYN, METOLAZONE

LANTHEUS MEDCL

* LANTHEUS MEDICAL IMAGING INC
 CARDIOLITE, TECHNETIUM TC-99M SESTAMIBI KIT
 DEFINITY, PERFLUTREN
 GALLIUM CITRATE GA 67, GALLIUM CITRATE GA-67
 NEUROLITE, TECHNETIUM TC-99M BICISATE KIT
 TECHNELITE, TECHNETIUM TC-99M SODIUM PERTECHNETATE GENERATOR
 THALLOUS CHLORIDE TL 201, THALLOUS CHLORIDE TL-201
 XENON XE 133, XENON XE-133

LANTHEUS MEDICAL

* LANTHEUS MEDICAL IMAGING INC
 FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** L ****

* LANTHEUS MEDICAL IMAGING INC
QUADRAMET, SAMARIUM SM-153 LEXIDRONAM PENTASODIUM

LARKEN LABS

* LARKEN LABORATORIES INC
DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE

LARKEN LABS INC

* LARKEN LABORATORIES INC
ACETAMINOPHEN, CAFFEINE AND DIHYDROCODEINE BITARTRATE, ACETAMINOPHEN
ALLZITAL, ACETAMINOPHEN
BUTALBITAL AND ACETAMINOPHEN, ACETAMINOPHEN
DEXAMETHASONE, DEXAMETHASONE

LAURUS LABS LTD

* LAURUS LABS LTD
METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE

LAVIPHARM LABS

* LAVIPHARM LABORATORIES INC
FENTANYL-100, FENTANYL
FENTANYL-25, FENTANYL
FENTANYL-50, FENTANYL
FENTANYL-75, FENTANYL

LEADIANT BIOSCI INC

* LEADIANT BIOSCIENCES INC
ABELCET, AMPHOTERICIN B
CARNITOR SF, LEVOCARNITINE
CARNITOR, LEVOCARNITINE
CYSTARAN, CYSTEAMINE HYDROCHLORIDE
MATULANE, PROCARBAZINE HYDROCHLORIDE

LEADING PHARMA LLC

* LEADING PHARMA LLC
BENZTROPINE MESYLATE, BENZTROPINE MESYLATE
CLOMIPRAMINE HYDROCHLORIDE, CLOMIPRAMINE HYDROCHLORIDE
DIPHENOXYLATE HYDROCHLORIDE AND ATROPINE SULFATE, ATROPINE SULFATE
FOLIC ACID, FOLIC ACID
FUROSEMIDE, FUROSEMIDE
GLYCOPYRROLATE, GLYCOPYRROLATE
HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
IMIPRAMINE HYDROCHLORIDE, IMIPRAMINE HYDROCHLORIDE
LORAZEPAM, LORAZEPAM
NIFEDIPINE, NIFEDIPINE

LEO LABS

* LEO LABORATORIES LTD
PICATO, INGENOL MEBUTATE

LEO PHARMA AS

* LEO PHARMA AS
DESONATE, DESONIDE
DOVONEX, CALCIPOTRIENE
ENSTILAR, BETAMETHASONE DIPROPIONATE
FINACEA, AZELAIC ACID
PROTOPIC, TACROLIMUS
TACLONEX, BETAMETHASONE DIPROPIONATE

LEXICON PHARMS INC

* LEXICON PHARMACEUTICALS INC
XERMELO, TELOTRISTAT ETIPRATE

LG CHEM LTD

* LG CHEM LTD
FACTIVE, GEMIFLOXACIN MESYLATE

LIEBEL-FLARSHEIM

* LIEBEL-FLARSHEIM CO LLC
CONRAY, IOTHALAMATE MEGLUMINE
CYSTO-CONRAY II, IOTHALAMATE MEGLUMINE
MD-GASTROVIEW, DIATRIZOATE MEGLUMINE
OPTIRAY 240, IOVERSOL
OPTIRAY 300, IOVERSOL
OPTIRAY 320, IOVERSOL

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** L ****

* LIEBEL-FLARSHEIM CO LLC
OPTIRAY 350, IOVERSOL
SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, SODIUM CHLORIDE

LIFE MOLECULAR

* LIFE MOLECULAR IMAGING SA
NEURACEQ, FLORBETABEN F-18

LIFEPHARMA

* LIFEPHARMA FZE
LACTULOSE, LACTULOSE

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)

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
LEVETIRACETAM, LEVETIRACETAM
* LOTUS PHARMACEUTICAL CO LTD NANTOU PLANT
LEVETIRACETAM, LEVETIRACETAM

LUITPOLD

* LUITPOLD PHARMACEUTICALS INC
AMINOCAPROIC ACID, AMINOCAPROIC ACID
BUSULFAN, BUSULFAN

LUKARE MEDICAL LLC

* LUKARE MEDICAL LLC
ELLIOTTS B SOLUTION, CALCIUM CHLORIDE

LUNDBECK NA LTD

* LUNDBECK NA LTD
NORTHERA, DROXIDOPA

LUNDBECK PHARMS LLC

* LUNDBECK PHARMACEUTICALS LLC
ONFI, CLOBAZAM
SABRIL, VIGABATRIN

LUPIN

* LUPIN INC
FLUOCINOLONE ACETONIDE, FLUOCINOLONE ACETONIDE
NALOXONE HYDROCHLORIDE AND PENTAZOCINE HYDROCHLORIDE, NALOXONE HYDROCHLORIDE
NYSTATIN, NYSTATIN
ORPHENADRINE CITRATE, ORPHENADRINE CITRATE
QUINARETIC, HYDROCHLOROTHIAZIDE
SOLOSEC, SECNIDAZOLE
TOBRAMYCIN, TOBRAMYCIN
TRIMETHOBENZAMIDE HYDROCHLORIDE, TRIMETHOBENZAMIDE HYDROCHLORIDE
* LUPIN LTD
AMLODIPINE BESYLATE AND VALSARTAN, AMLODIPINE BESYLATE
AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
CARVEDILOL, CARVEDILOL
CEFADROXIL, CEFADROXIL/CEFADROXIL HEMIHYDRATE
CEFDINIR, CEFDINIR
CEFPROZIL, CEFPROZIL
CEFTRIAZONE, CEFTRIAZONE SODIUM
CEFUROXIME AXETIL, CEFUROXIME AXETIL
CEPHALEXIN, CEPHALEXIN
DIVALPROEX SODIUM, DIVALPROEX SODIUM
ETHAMBUTOL HYDROCHLORIDE, ETHAMBUTOL HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** L ******* LUPIN LTD**

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 ATLANTIS*** LUPIN ATLANTIS HOLDINGS SA**

ANTARA (MICRONIZED), FENOFIBRATE
 BUDESONIDE, BUDESONIDE
 DESOXIMETASONE, DESOXIMETASONE
 LEVOTHYROXINE SODIUM, LEVOTHYROXINE SODIUM **
 MIBELAS 24 FE, ETHINYL ESTRADIOL
 OSELTAMIVIR PHOSPHATE, OSELTAMIVIR PHOSPHATE
 TESTOSTERONE, TESTOSTERONE
 TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE

LUPIN LTD*** LUPIN LIMITED**

LEVETIRACETAM, LEVETIRACETAM
 LEVONORGESTREL AND ETHINYL ESTRADIOL AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL

*** LUPIN LTD**

ABACAVIR SULFATE AND LAMIVUDINE, ABACAVIR SULFATE
 ABACAVIR SULFATE, LAMIVUDINE AND ZIDOVUDINE, ABACAVIR SULFATE
 AMABELZ, ESTRADIOL
 AMLODIPINE BESYLATE, VALSARTAN AND HYDROCHLOROTHIAZIDE, AMLODIPINE BESYLATE
 ARMODAFINIL, ARMODAFINIL
 ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM
 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
 BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
 CALCIUM ACETATE, CALCIUM ACETATE
 CELECOXIB, CELECOXIB
 CLOBAZAM, CLOBAZAM
 CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
 CLOMIPRAMINE HYDROCHLORIDE, CLOMIPRAMINE HYDROCHLORIDE
 CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
 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
 ENSKYCE, DESOGESTREL
 ESCITALOPRAM OXALATE, ESCITALOPRAM OXALATE
 ESZOPICLONE, ESZOPICLONE
 ETHACRYNIC ACID, ETHACRYNIC ACID
 EXTENDED PHENYTOIN SODIUM, PHENYTOIN SODIUM
 FALLBACK SOLO, LEVONORGESTREL (OTC)

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** L **

* LUPIN LTD
 FAMOTIDINE, FAMOTIDINE
 FAYOSIM, ETHINYL ESTRADIOL
 FENOFIBRATE, FENOFIBRATE
 FENOFIBRIC ACID, CHOLINE FENOFIBRATE
 FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 FOSAPREPITANT DIMEGLUMINE, FOSAPREPITANT DIMEGLUMINE
 FYAVOLV, ETHINYL ESTRADIOL
 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
 MEFENAMIC ACID, MEFENAMIC ACID
 MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 METRONIDAZOLE, METRONIDAZOLE
 MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE
 MOXIFLOXACIN HYDROCHLORIDE, MOXIFLOXACIN HYDROCHLORIDE
 NABUMETONE, NABUMETONE
 NADOLOL, NADOLOL
 NIACIN, NIACIN
 NIKKI, DROSPIRENONE
 NORETHINDRONE AND ETHINYL ESTRADIOL AND FERROUS FUMARATE, ETHINYL ESTRADIOL
 NORETHINDRONE, NORETHINDRONE
 NORGESTIMATE AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
 NYSTATIN AND TRIAMCINOLONE ACETONIDE, NYSTATIN
 OLMESARTAN MEDOXOMIL, OLMESARTAN MEDOXOMIL
 PAROXETINE HYDROCHLORIDE, PAROXETINE HYDROCHLORIDE
 PIOGLITAZONE HYDROCHLORIDE, PIOGLITAZONE HYDROCHLORIDE
 PIRMELLA 1/35, ETHINYL ESTRADIOL
 PIRMELLA 7/7/7, ETHINYL ESTRADIOL
 POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
 QUETIAPINE FUMARATE, QUETIAPINE FUMARATE
 QUININE SULFATE, QUININE SULFATE
 RABEPRAZOLE SODIUM, RABEPRAZOLE SODIUM
 RANOLAZINE, RANOLAZINE
 RIFABUTIN, RIFABUTIN
 ROSUVASTATIN CALCIUM, ROSUVASTATIN CALCIUM
 SILDENAFIL CITRATE, SILDENAFIL CITRATE
 SILODOSIN, SILODOSIN
 SUPRAX, CEFIXIME
 TADALAFIL, TADALAFIL
 TELMISARTAN AND AMLODIPINE, AMLODIPINE BESYLATE
 TELMISARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 TESTOSTERONE, TESTOSTERONE
 TETRABENAZINE, TETRABENAZINE
 TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE
 TYDEMY, DROSPIRENONE
 VALSARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 VALSARTAN, VALSARTAN

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** L ****

* LUPIN LTD
 VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE
 VYFEMLA, ETHINYL ESTRADIOL
 ZILEUTON, ZILEUTON
 ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE

LUPIN PHARMS

* LUPIN PHARMACEUTICALS INC
 AMLODIPINE BESYLATE AND BENAZEPRIL HYDROCHLORIDE, AMLODIPINE BESYLATE
 DESLORATADINE, DESLORATADINE
 MELOXICAM, MELOXICAM
 NORGESTIMATE AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
 PRAVASTATIN SODIUM, PRAVASTATIN SODIUM
 RIFAMPIN, RIFAMPIN
 SUPRAX, CEFIXIME
 ZIPRASIDONE HYDROCHLORIDE, ZIPRASIDONE HYDROCHLORIDE

LYMOL MEDCL

* LYMOL MEDICAL CORP
 SCLEROSOL, TALC
 TALC, TALC

LYNE

* LYNE LABORATORIES INC
 CHLORHEXIDINE GLUCONATE, CHLORHEXIDINE GLUCONATE
 CLINDAMYCIN PALMITATE HYDROCHLORIDE, CLINDAMYCIN PALMITATE HYDROCHLORIDE
 CYPROHEPTADINE HYDROCHLORIDE, CYPROHEPTADINE HYDROCHLORIDE
 DEXAMETHASONE, DEXAMETHASONE
 ERYTHROMYCIN AND BENZOYL PEROXIDE, BENZOYL PEROXIDE
 FLUOCINOLONE ACETONIDE, FLUOCINOLONE ACETONIDE
 LEVOCARNITINE, LEVOCARNITINE
 NYSTATIN, NYSTATIN
 TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE

PERRIGO

* L PERRIGO CO
 ACETAMINOPHEN, ACETAMINOPHEN (OTC)
 ACETAMINOPHEN, ASPIRIN AND CAFFEINE, ACETAMINOPHEN (OTC)
 CHILDREN'S IBUPROFEN, IBUPROFEN (OTC)
 CROMOLYN SODIUM, CROMOLYN SODIUM (OTC)
 DOXYLAMINE SUCCINATE, DOXYLAMINE SUCCINATE (OTC)
 FAMOTIDINE, FAMOTIDINE (OTC)
 IBUPROFEN AND PSEUDOEPHEDRINE HYDROCHLORIDE, IBUPROFEN (OTC)
 IBUPROFEN, IBUPROFEN (OTC)
 LOPERAMIDE HYDROCHLORIDE AND SIMETHICONE, LOPERAMIDE HYDROCHLORIDE (OTC)
 LOPERAMIDE HYDROCHLORIDE, LOPERAMIDE HYDROCHLORIDE (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)
 RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE (OTC)
 TAB-PROFEN, IBUPROFEN (OTC)
 TIOCONAZOLE, TIOCONAZOLE (OTC)

**** M ******MA GENERAL HOSP**

* MASSACHUSETTS GENERAL HOSP
 FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18

MACLEODS PHARMS LTD

* MACLEODS PHARMACEUTICALS LTD
 AMLODIPINE AND OLMESARTAN MEDOXOMIL, AMLODIPINE BESYLATE
 AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
 ARIPIPIRAZOLE, ARIPIPIRAZOLE
 CANDESARTAN CILEXETIL AND HYDROCHLOROTHIAZIDE, CANDESARTAN CILEXETIL
 CANDESARTAN CILEXETIL, CANDESARTAN CILEXETIL
 CELECOXIB, CELECOXIB
 CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** M ****

* MACLEODS PHARMACEUTICALS LTD
 CLOPIDOGREL BISULFATE, CLOPIDOGREL BISULFATE
 DARIFENACIN, DARIFENACIN HYDROBROMIDE
 DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
 DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE
 ENTACAPONE, ENTACAPONE
 ESCITALOPRAM OXALATE, ESCITALOPRAM OXALATE
 ESZOPICLONE, ESZOPICLONE
 FAMCICLOVIR, FAMCICLOVIR
 FLUOCINONIDE, FLUOCINONIDE
 IBANDRONATE SODIUM, IBANDRONATE SODIUM
 IRBESARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 IRBESARTAN, IRBESARTAN
 LAMIVUDINE AND ZIDOVUDINE, LAMIVUDINE
 LAMIVUDINE, LAMIVUDINE
 LEVOCETIRIZINE DIHYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE
 LEVOFLOXACIN, LEVOFLOXACIN
 LIDOCAINE, LIDOCAINE
 LOSARTAN POTASSIUM AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 LOSARTAN POTASSIUM, LOSARTAN POTASSIUM
 MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 MONTELUKAST SODIUM, MONTELUKAST SODIUM
 NEVIRAPINE, NEVIRAPINE
 OLANZAPINE, OLANZAPINE
 OLMESARTAN MEDOXOMIL, OLMESARTAN MEDOXOMIL
 OSELTAMIVIR PHOSPHATE, OSELTAMIVIR PHOSPHATE
 PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
 PIOGLITAZONE HYDROCHLORIDE AND METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 PIOGLITAZONE HYDROCHLORIDE, PIOGLITAZONE HYDROCHLORIDE
 PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE
 QUETIAPINE FUMARATE, QUETIAPINE FUMARATE
 RISEDRONATE SODIUM, RISEDRONATE SODIUM
 RIVASTIGMINE TARTRATE, RIVASTIGMINE TARTRATE
 RIZATRIPTAN BENZOATE, RIZATRIPTAN BENZOATE
 SILDENAFIL CITRATE, SILDENAFIL CITRATE
 SILODOSIN, SILODOSIN
 TADALAFIL, TADALAFIL
 TAMSULOSIN HYDROCHLORIDE, TAMSULOSIN HYDROCHLORIDE
 TELMISARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 TENOFOVIR DISOPROXIL FUMARATE, TENOFOVIR DISOPROXIL FUMARATE
 TOLTERODINE TARTRATE, TOLTERODINE TARTRATE
 TRAMADOL HYDROCHLORIDE AND ACETAMINOPHEN, ACETAMINOPHEN
 TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE
 TRIAMCINOLONE ACETATE, TRIAMCINOLONE ACETONIDE
 TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE
 VALSARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 VALSARTAN, VALSARTAN
 VARDENAFIL HYDROCHLORIDE, VARDENAFIL HYDROCHLORIDE
 VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE
 ZIPRASIDONE HYDROCHLORIDE, ZIPRASIDONE HYDROCHLORIDE

MAIA PHARMS INC

* MAIA PHARMACEUTICALS INC
 ANGIOMAX RTU, BIVALIRUDIN
 BACLOFEN, BACLOFEN
 LEVOTHYROXINE SODIUM, LEVOTHYROXINE SODIUM
 SODIUM PHENYLACETATE AND SODIUM BENZOATE, SODIUM BENZOATE

MAINPOINTE

* MAINPOINTE PHARMACEUTICALS LLC
 TUXARIN ER, CHLORPHENIRAMINE MALEATE

MALLINCKRODT ARD

* MALLINCKRODT ARD INC
 H.P. ACTHAR GEL, CORTICOTROPIN

MALLINCKRODT HOSP

* MALLINCKRODT HOSP PRODUCTS IP LTD

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** M ****

* MALLINCKRODT HOSP PRODUCTS IP LTD
 INOMAX, NITRIC OXIDE
 OFIRMEV, ACETAMINOPHEN
 UVADEX, METHOXSALEN

MANKIND PHARMA

* MANKIND PHARMA LTD
 CLOMIPRAMINE HYDROCHLORIDE, CLOMIPRAMINE HYDROCHLORIDE
 RANOLAZINE, RANOLAZINE

MANNKIND

* MANNKIND CORP
 AFREZZA, INSULIN RECOMBINANT HUMAN

MARKSANS PHARMA

* MARKSANS PHARMA LTD
 CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 CETIRIZINE HYDROCHLORIDE HIVES, CETIRIZINE HYDROCHLORIDE (OTC)
 DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE
 DUTASTERIDE, DUTASTERIDE
 FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 GABAPENTIN, GABAPENTIN
 IBUPROFEN, IBUPROFEN
 IBUPROFEN, IBUPROFEN (OTC)
 LORATADINE, LORATADINE (OTC)
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 NAPROXEN SODIUM, NAPROXEN SODIUM (OTC)
 NAPROXEN, NAPROXEN
 PARICALCITOL, PARICALCITOL

MARNEL PHARMS

* MARNEL PHARMACEUTICALS LLC
 ALA-SCALP, HYDROCORTISONE
 CROTAN, CROTAMITON

MAYER LABS INC

* MAYER LABORATORIES INC
 TODAY, NONOXYNOL-9 (OTC)

MAYNE PHARMA

* MAYNE PHARMA INTERNATIONAL PTY LTD
 DORYX MPC, DOXYCYCLINE HYCLATE
 DORYX, DOXYCYCLINE HYCLATE
 ERYC, ERYTHROMYCIN

* MAYNE PHARMA LLC
 BUDESONIDE, BUDESONIDE
 CAMILA, NORETHINDRONE
 CARBIDOPA AND LEVODOPA, CARBIDOPA
 CLARITHROMYCIN, CLARITHROMYCIN
 CLONIDINE, CLONIDINE
 CLOZAPINE, CLOZAPINE
 CYCLOSPORINE, CYCLOSPORINE
 DESOGESTREL AND ETHINYL ESTRADIOL, DESOGESTREL
 DEXTROAMPHETAMINE SULFATE, DEXTROAMPHETAMINE SULFATE
 DIAZEPAM, DIAZEPAM
 DISOPYRAMIDE PHOSPHATE, DISOPYRAMIDE PHOSPHATE
 DROSPIRENONE AND ETHINYL ESTRADIOL, DROSPIRENONE
 ERRIN, NORETHINDRONE
 ESTAZOLAM, ESTAZOLAM
 ESTRADIOL, ESTRADIOL
 FABIOR, TAZAROTENE
 FLUOROURACIL, FLUOROURACIL
 LEVONORGESTREL AND ETHINYL ESTRADIOL AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
 LEVONORGESTREL AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
 LEVORA 0.15/30-28, ETHINYL ESTRADIOL
 LEXETTE, HALOBETASOL PROPIONATE
 LOW-OGESTREL-28, ETHINYL ESTRADIOL
 METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
 MICROGESTIN 1.5/30, ETHINYL ESTRADIOL
 MICROGESTIN 1/20, ETHINYL ESTRADIOL
 MICROGESTIN FE 1.5/30, ETHINYL ESTRADIOL

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** M ******* MAYNE PHARMA LLC**

MICROGESTIN FE 1/20, ETHINYL ESTRADIOL
 NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
 NORETHINDRONE AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
 NORTRIPTYLINE HYDROCHLORIDE, NORTRIPTYLINE HYDROCHLORIDE
 SORILUX, CALCIPOTRIENE
 TAMOXIFEN CITRATE, TAMOXIFEN CITRATE
 TOLCAPONE, TOLCAPONE
 TRI-NORINYL 28-DAY, ETHINYL ESTRADIOL
 TRIMETHOPRIM, TRIMETHOPRIM
 TRIVORA-28, ETHINYL ESTRADIOL
 ZOVIA 1/35E-28, ETHINYL ESTRADIOL

MAYNE PHARMA INC*** MAYNE PHARMA INC**

AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE
 BROMPHENIRAMINE MALEATE, PSEUDOEPHEDRINE HYDROCHLORIDE AND DEXTROMETHORPHAN
 BUTALBITAL AND ACETAMINOPHEN, ACETAMINOPHEN
 BUTALBITAL, ACETAMINOPHEN AND CAFFEINE, ACETAMINOPHEN
 BUTALBITAL, ASPIRIN, CAFFEINE, AND CODEINE PHOSPHATE, ASPIRIN
 CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
 DOFETILIDE, DOFETILIDE
 DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
 DOXYCYCLINE, DOXYCYCLINE
 LIOTHYRONINE SODIUM, LIOTHYRONINE SODIUM
 METHAMPHETAMINE HYDROCHLORIDE, METHAMPHETAMINE HYDROCHLORIDE
 MORPHINE SULFATE, MORPHINE SULFATE
 NYSTATIN, NYSTATIN
 OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN
 OXYCODONE AND ASPIRIN, ASPIRIN
 OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE

MAYNE PHARMA INTL*** MAYNE PHARMA INTERNATIONAL PTY LTD**

TOLSURA, ITRACONAZOLE

MCGUFF*** MCGUFF PHARMACEUTICALS INC**

ASCOR, ASCORBIC ACID

MCNEIL*** MCNEIL CONSUMER PRODUCTS CO DIV MCNEILAB INC**

IBUPROFEN, IBUPROFEN (OTC)

MCNEIL CONS*** MCNEIL CONSUMER HEALTHCARE**

SUDAFED 12 HOUR, PSEUDOEPHEDRINE HYDROCHLORIDE (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

MDGH*** MEDICINES DEVELOPMENT FOR GLOBAL HEALTH**

MOXIDECTIN, MOXIDECTIN

MEDEFIL INC*** MEDEFIL INC**

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

MEDICINES360*** MEDICINES360**

LILETTA, LEVONORGESTREL

MEDICIS*** MEDICIS PHARMACEUTICAL CORP**

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** M ****

* MEDICIS PHARMACEUTICAL CORP
 AMMONUL, SODIUM BENZOATE
 CALCIUM DISODIUM VERSENATE, EDETATE CALCIUM DISODIUM
 LUZU, LULICONAZOLE
 SOLODYN, MINOCYCLINE HYDROCHLORIDE
 VANOS, FLUOCINONIDE
 ZIANA, CLINDAMYCIN PHOSPHATE

MEDICURE

* MEDICURE INTERNATIONAL INC
 AGGRASTAT, TIROFIBAN HYDROCHLORIDE
 SODIUM NITROPRUSSIDE, SODIUM NITROPRUSSIDE
 ZYPITAMAG, PITAVASTATIN MAGNESIUM

MEDIMETRIKS PHARMS

* MEDIMETRIKS PHARMACEUTICALS INC
 BETAXOLOL HYDROCHLORIDE, BETAXOLOL HYDROCHLORIDE
 LOPROX, CICLOPIROX
 NEO-SYNALAR, FLUOCINOLONE ACETONIDE
 SYNALAR, FLUOCINOLONE ACETONIDE

MEDLINE INDUSTRIES

* MEDLINE INDUSTRIES INC
 READYPREP CHG, CHLORHEXIDINE GLUCONATE (OTC)

MEDTECH PRODUCTS

* MEDTECH 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)

MEITHEAL

* MEITHEAL PHARMACEUTICALS INC
 ATRACURIUM BESYLATE PRESERVATIVE FREE, ATRACURIUM BESYLATE
 ATRACURIUM BESYLATE, ATRACURIUM BESYLATE
 BLEOMYCIN SULFATE, BLEOMYCIN SULFATE
 CARBOPLATIN, CARBOPLATIN
 CISATRACURIUM BESYLATE, CISATRACURIUM BESYLATE
 CYTARABINE, CYTARABINE
 DOXERCALCIFEROL, DOXERCALCIFEROL
 LEVOLEUCOVORIN CALCIUM, LEVOLEUCOVORIN CALCIUM
 PHENYLEPHRINE HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE
 TOPOTECAN HYDROCHLORIDE, TOPOTECAN HYDROCHLORIDE

MELINTA

* MELINTA SUBSIDIARY CORP
 BAXDELA, DELAFLOXACIN MEGLUMINE

MELINTA THERAP

* MELINTA THERAPEUTICS INC
 ORBACTIV, ORITAVANCIN DIPHOSPHATE

MEM SLOAN-KETTERING

* MEMORIAL SLOAN-KETTERING CANCER CENTER
 FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18

MERCK

* MERCK AND CO INC
 CANCIDAS, CASPOFUNGIN ACETATE
 EMEND, APREPITANT
 FOSAMAX PLUS D, ALENDRONATE SODIUM
 MAXALT, RIZATRIPTAN BENZOATE
 MAXALT-MLT, RIZATRIPTAN BENZOATE
 PRIMAXIN, CILASTATIN SODIUM
 PROSCAR, FINASTERIDE
 ZOLINZA, VORINOSTAT
 * MERCK RESEARCH LABORATORIES DIV MERCK CO INC
 PRINIVIL, LISINOPRIL

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** M ****

* MERCK RESEARCH LABORATORIES DIV MERCK CO INC
 PROPECIA, FINASTERIDE
 SINGULAIR, MONTELUKAST SODIUM
 TRUSOPT, DORZOLAMIDE HYDROCHLORIDE

MERCK AND CO INC

* MERCK AND CO INC
 EMEND, FOSAPREPITANT DIMEGLUMINE
 FOSAMAX, ALENDRONATE SODIUM

MERCK SHARP DOHME

* MERCK SHARP AND DOHME CORP
 ASMANEX HFA, MOMETASONE FUROATE
 ASMANEX TWISTHALER, MOMETASONE FUROATE
 BELSOMRA, SUVOREXANT
 CELESTONE SOLUSPAN, BETAMETHASONE ACETATE
 CLARINEX D 24 HOUR, DESLORATADINE
 CLARINEX, DESLORATADINE
 CLARINEX-D 12 HOUR, DESLORATADINE
 COZAAR, LOSARTAN POTASSIUM
 CRIXIVAN, INDINAVIR SULFATE
 DIPROLENE, BETAMETHASONE DIPROPIONATE
 DULERA, FORMOTEROL FUMARATE
 ELOCON, MOMETASONE FUROATE
 GUANIDINE HYDROCHLORIDE, GUANIDINE HYDROCHLORIDE
 HYZAAR, HYDROCHLOROTHIAZIDE
 INVANZ, ERTAPENEM SODIUM
 ISENTRESS HD, RALTEGRAVIR POTASSIUM
 ISENTRESS, RALTEGRAVIR POTASSIUM
 JANUMET XR, METFORMIN HYDROCHLORIDE
 JANUMET, METFORMIN HYDROCHLORIDE
 JANUVIA, SITAGLIPTIN PHOSPHATE
 LOTRISONE, BETAMETHASONE DIPROPIONATE
 NASONEX, MOMETASONE FUROATE
 NOXAFIL, POSACONAZOLE
 PREVYMIS, LETERMIVIR
 REBETOL, RIBAVIRIN
 SEGLUROMET, ERTUGLIFLOZIN
 SINEMET, CARBIDOPA
 STEGLATRO, ERTUGLIFLOZIN
 STEGLUJAN, ERTUGLIFLOZIN
 STROMECTOL, IVERMECTIN
 TEMODAR, TEMOZOLOMIDE
 ZEPATIER, ELBASVIR

MERIDIAN MEDCL

* MERIDIAN MEDICAL TECHNOLOGIES INC
 DUODOTE, ATROPINE

MERIDIAN MEDCL TECHN

* MERIDIAN MEDICAL TECHNOLOGIES INC
 ATROPEN, ATROPINE
 MORPHINE SULFATE, MORPHINE SULFATE
 PRALIDOXIME CHLORIDE, PRALIDOXIME CHLORIDE
 SEIZALAM, MIDAZOLAM HYDROCHLORIDE

MERRO PHARM

* MERRO PHARMACEUTICAL CO LTD
 IBUPROFEN, IBUPROFEN (OTC)

MERZ PHARMS

* MERZ PHARMACEUTICALS LLC
 CUVPOSA, GLYCOPYRROLATE

METACEL PHARMS LLC

* METACEL PHARMACEUTICALS LLC
 OZOBAX, BACLOFEN

METHAPHARM

* METHAPHARM INC
 PROVOCHOLINE, METHACHOLINE CHLORIDE

METUCHEN PHARMS

* METUCHEN PHARMACEUTICALS LLC

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** M ****

* METUCHEN PHARMACEUTICALS LLC
STENDRA, AVANAFIL

MICRO LABS

* MICRO LABS LTD
AMLODIPINE AND OLMESARTAN MEDOXOMIL, AMLODIPINE BESYLATE
APIXABAN, APIXABAN
CAFFEINE CITRATE, CAFFEINE CITRATE
CELECOXIB, CELECOXIB
CLINDAMYCIN HYDROCHLORIDE, CLINDAMYCIN HYDROCHLORIDE
CLOBAZAM, CLOBAZAM
DALFAMPRIDINE, DALFAMPRIDINE
DIPHENHYDRAMINE HYDROCHLORIDE, DIPHENHYDRAMINE HYDROCHLORIDE
DORZOLAMIDE HYDROCHLORIDE, DORZOLAMIDE HYDROCHLORIDE
GLIMEPIRIDE, GLIMEPIRIDE
LEVETIRACETAM, LEVETIRACETAM
LEVOCETIRIZINE DIHYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE (OTC)
LOSARTAN POTASSIUM, LOSARTAN POTASSIUM
MEFENAMIC ACID, MEFENAMIC ACID
METHENAMINE HIPPURATE, METHENAMINE HIPPURATE
OLMESARTAN MEDOXOMIL, OLMESARTAN MEDOXOMIL
PIROXICAM, PIROXICAM
RANOLAZINE, RANOLAZINE
RASAGILINE MESYLATE, RASAGILINE MESYLATE
ROFLUMILAST, ROFLUMILAST
SIMVASTATIN, SIMVASTATIN
SODIUM NITROPRUSSIDE, SODIUM NITROPRUSSIDE
TAFLUPROST, TAFLUPROST
TRANEXAMIC ACID, TRANEXAMIC ACID
VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE

MICRO LABS LTD

* MICRO LABS LTD
NEVIRAPINE, NEVIRAPINE

MICRO LABS LTD INDIA

* MICRO LABS LTD INDIA
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

MIDATECH PHARMA US

* MIDATECH PHARMA US INC
ZUPLENZ, ONDANSETRON

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
BENZONATATE, BENZONATATE
BUTALBITAL AND ACETAMINOPHEN, ACETAMINOPHEN
BUTALBITAL, ACETAMINOPHEN AND CAFFEINE, ACETAMINOPHEN
BUTAPAP, ACETAMINOPHEN
CARBINOXAMINE MALEATE, CARBINOXAMINE MALEATE
CHLORZOAZONE, CHLORZOAZONE
ERGOTAMINE TARTRATE AND CAFFEINE, CAFFEINE
ETHOSUXIMIDE, ETHOSUXIMIDE
HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
METHAZOLAMIDE, METHAZOLAMIDE
OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN
PHENDIMETRAZINE TARTRATE, PHENDIMETRAZINE TARTRATE
TRIHENYPHENIDYL HYDROCHLORIDE, TRIHENYPHENIDYL HYDROCHLORIDE

MIKART INC

* MIKART INC

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** M ****

* MIKART INC
OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN

MILLENNIUM PHARMS

* MILLENNIUM PHARMACEUTICALS INC
NINLARO, IXAZOMIB CITRATE
VELCADE, BORTEZOMIB

MILLICENT

* MILLICENT HOLDINGS LTD
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

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 CO

* MISSION PHARMACAL CO
CARBINOXAMINE MALEATE, CARBINOXAMINE MALEATE
POTASSIUM IODIDE, POTASSIUM IODIDE (OTC)
THIOLA EC, TIOPRONIN

MITSUBISHI TANABE

* MITSUBISHI TANABE PHARMA CORP
RADICAVA, EDARAVONE

MLV

* MLV PHARMA LLC
ACETIC ACID, ACETIC ACID, GLACIAL
CETIRIZINE HYDROCHLORIDE, CETIRIZINE HYDROCHLORIDE
ROPINIROLE HYDROCHLORIDE, ROPINIROLE HYDROCHLORIDE
TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE

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
CORTISPORIN, BACITRACIN ZINC
CORTISPORIN, HYDROCORTISONE ACETATE
MENEST, ESTROGENS, ESTERIFIED
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
GRISEOFULVIN, ULTRAMICROSIZE, GRISEOFULVIN, ULTRAMICROSIZE
METAXALONE, METAXALONE
METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
POTASSIUM CITRATE, POTASSIUM CITRATE

MSD INTL

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** M ****

* MSD INTERNATIONAL GMBH
VYTORIN, EZETIMIBE

MSD INTL GMBH

* MSD INTERNATIONAL GMBH
ZETIA, EZETIMIBE

MSD MERCK CO

* MERCK SHARP AND DOHME CORP A SUB OF MERCK AND CO INC
DELSTRIGO, DORAVIRINE
EMEND, APREPITANT
PIFELTRO, DORAVIRINE
RECARBRIO, CILASTATIN SODIUM
SINGULAIR, MONTELUKAST SODIUM
ZOCOR, SIMVASTATIN

MSN

* MSN LABORATORIES PRIVATE LTD
ABIRATERONE ACETATE, ABIRATERONE ACETATE
ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM
CAPECITABINE, CAPECITABINE
CLOFARABINE, CLOFARABINE
DECITABINE, DECITABINE
DEFERASIROX, DEFERASIROX
DOFETILIDE, DOFETILIDE
FEBUXOSTAT, FEBUXOSTAT
FOSAPREPITANT DIMEGLUMINE, FOSAPREPITANT DIMEGLUMINE
MOXIFLOXACIN HYDROCHLORIDE, MOXIFLOXACIN HYDROCHLORIDE
PREGABALIN, PREGABALIN
ROSUVASTATIN CALCIUM, ROSUVASTATIN CALCIUM
SILODOSIN, SILODOSIN
SOLIFENACIN SUCCINATE, SOLIFENACIN SUCCINATE
THIOTEPA, THIOTEPA
TRIENTINE HYDROCHLORIDE, TRIENTINE HYDROCHLORIDE

MURTY PHARMS

* MURTY PHARMACEUTICALS INC
AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE
DIPYRIDAMOLE, DIPYRIDAMOLE

MYLAN

* MYLAN PHARMACEUTICALS
FENOFIBRATE, FENOFIBRATE
METOPROLOL TARTRATE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
TRANEXAMIC ACID, TRANEXAMIC ACID

* MYLAN PHARMACEUTICALS INC
ABIRATERONE ACETATE, ABIRATERONE ACETATE
ACAMPROSATE CALCIUM, ACAMPROSATE CALCIUM
ACEBUTOLOL HYDROCHLORIDE, ACEBUTOLOL HYDROCHLORIDE
ACITRETIN, ACITRETIN
ALBUTEROL SULFATE, ALBUTEROL SULFATE
ALLOPURINOL, ALLOPURINOL
ALMOTRIPTAN MALATE, ALMOTRIPTAN MALATE
ALPRAZOLAM, ALPRAZOLAM
AMBRISENTAN, AMBRISENTAN
AMILORIDE HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, AMILORIDE HYDROCHLORIDE
AMITRIPTYLINE HYDROCHLORIDE, AMITRIPTYLINE HYDROCHLORIDE
AMLODIPINE BESYLATE AND ATORVASTATIN CALCIUM, AMLODIPINE BESYLATE
AMLODIPINE BESYLATE AND VALSARTAN, AMLODIPINE BESYLATE
AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
ATENOLOL AND CHLORTHALIDONE, ATENOLOL
ATENOLOL, ATENOLOL
ATOVAQUONE AND PROGUANIL HYDROCHLORIDE, ATOVAQUONE
AVITA, TRETINOIN
AZATHIOPRINE, AZATHIOPRINE
BENAZEPRIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, BENAZEPRIL HYDROCHLORIDE
BENAZEPRIL HYDROCHLORIDE, BENAZEPRIL HYDROCHLORIDE
BROMFENAC SODIUM, BROMFENAC SODIUM
BROMOCRIPTINE MESYLATE, BROMOCRIPTINE MESYLATE
BUDESONIDE, BUDESONIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** M **

* MYLAN PHARMACEUTICALS INC
 BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
 BUSPIRONE HYDROCHLORIDE, BUSPIRONE HYDROCHLORIDE
 BUTORPHANOL TARTRATE, BUTORPHANOL TARTRATE
 CANDESARTAN CILEXETIL AND HYDROCHLOROTHIAZIDE, CANDESARTAN CILEXETIL
 CANDESARTAN CILEXETIL, CANDESARTAN CILEXETIL
 CAPECITABINE, CAPECITABINE
 CAPTOPRIL AND HYDROCHLOROTHIAZIDE, CAPTOPRIL
 CAPTOPRIL, CAPTOPRIL
 CARBIDOPA AND LEVODOPA, CARBIDOPA
 CARVEDILOL, CARVEDILOL
 CELECOXIB, CELECOXIB
 CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 CETIRIZINE HYDROCHLORIDE HIVES, CETIRIZINE HYDROCHLORIDE (OTC)
 CHLOROTHIAZIDE, CHLOROTHIAZIDE
 CHLORTHALIDONE, CHLORTHALIDONE
 CIMETIDINE, CIMETIDINE
 CINACALCET HYDROCHLORIDE, CINACALCET HYDROCHLORIDE
 CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE
 CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE
 CLOBAZAM, CLOBAZAM
 CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
 CLOMIPRAMINE HYDROCHLORIDE, CLOMIPRAMINE HYDROCHLORIDE
 CLONAZEPAM, CLONAZEPAM
 CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
 CLORAZEPATE DIPOTASSIUM, CLORAZEPATE DIPOTASSIUM
 CLOZAPINE, CLOZAPINE
 COLCHICINE, COLCHICINE
 CYSTAGON, CYSTEAMINE BITARTRATE
 DENAVIR, PENCICLOVIR
 DESMOPRESSIN ACETATE, DESMOPRESSIN ACETATE
 DEXMETHYLPHENIDATE HYDROCHLORIDE, DEXMETHYLPHENIDATE HYDROCHLORIDE
 DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE
 DIAZEPAM, DIAZEPAM
 DICLOFENAC POTASSIUM, DICLOFENAC POTASSIUM
 DICLOFENAC SODIUM, DICLOFENAC SODIUM
 DICYCLOMINE HYDROCHLORIDE, DICYCLOMINE HYDROCHLORIDE
 DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
 DIPHENOXYLATE HYDROCHLORIDE AND ATROPINE SULFATE, ATROPINE SULFATE
 DIVALPROEX SODIUM, DIVALPROEX SODIUM
 DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
 DOXAZOSIN MESYLATE, DOXAZOSIN MESYLATE
 DOXYCYCLINE HCLATE, DOXYCYCLINE HCLATE
 ECONAZOLE NITRATE, ECONAZOLE NITRATE
 EFAVIRENZ, EFAVIRENZ
 ELETRIPTAN HYDROBROMIDE, ELETRIPTAN HYDROBROMIDE
 ENALAPRIL MALEATE AND HYDROCHLOROTHIAZIDE, ENALAPRIL MALEATE
 EPLERENONE, EPLERENONE
 ERLOTINIB HYDROCHLORIDE, ERLOTINIB HYDROCHLORIDE
 ERYGEL, ERYTHROMYCIN
 ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM
 ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM (OTC)
 ESTRADIOL, ESTRADIOL
 ETOPOSIDE, ETOPOSIDE
 EVOCLIN, CLINDAMYCIN PHOSPHATE
 EXEMESTANE, EXEMESTANE
 EXTENDED PHENYTOIN SODIUM, PHENYTOIN SODIUM
 EXTINA, KETOCONAZOLE
 FAMCICLOVIR, FAMCICLOVIR
 FEBUXOSTAT, FEBUXOSTAT
 FENOFIBRATE, FENOFIBRATE
 FEXOFENADINE HYDROCHLORIDE ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
 FEXOFENADINE HYDROCHLORIDE HIVES, FEXOFENADINE HYDROCHLORIDE (OTC)
 FEXOFENADINE HYDROCHLORIDE, FEXOFENADINE HYDROCHLORIDE
 FLUOROURACIL, FLUOROURACIL

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** M **

* MYLAN PHARMACEUTICALS INC
 FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 FLUPHENAZINE HYDROCHLORIDE, FLUPHENAZINE HYDROCHLORIDE
 FLURBIPROFEN, FLURBIPROFEN
 FOSAMPRENAVIR CALCIUM, FOSAMPRENAVIR CALCIUM
 FROVATRIPTAN SUCCINATE, FROVATRIPTAN SUCCINATE
 FUROSEMIDE, FUROSEMIDE
 GABAPENTIN, GABAPENTIN
 GATIFLOXACIN, GATIFLOXACIN
 GLATIRAMER ACETATE, GLATIRAMER ACETATE
 GLIMEPIRIDE, GLIMEPIRIDE
 GLIPIZIDE, GLIPIZIDE
 GUANFACINE HYDROCHLORIDE, GUANFACINE HYDROCHLORIDE
 HALCINONIDE, HALCINONIDE
 HALOPERIDOL, HALOPERIDOL
 HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 HYDROXYCHLOROQUINE SULFATE, HYDROXYCHLOROQUINE SULFATE
 IMATINIB MESYLATE, IMATINIB MESYLATE
 INDAPAMIDE, INDAPAMIDE
 INDOMETHACIN, INDOMETHACIN
 KETOCONAZOLE, KETOCONAZOLE
 KETOPROFEN, KETOPROFEN
 KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
 LAMOTRIGINE, LAMOTRIGINE
 LANSOPRAZOLE, LANSOPRAZOLE
 LANSOPRAZOLE, LANSOPRAZOLE (OTC)
 LEVETIRACETAM, LEVETIRACETAM
 LEVOTHYROXINE SODIUM, LEVOTHYROXINE SODIUM **
 LOPERAMIDE HYDROCHLORIDE, LOPERAMIDE HYDROCHLORIDE
 LORATADINE, LORATADINE (OTC)
 LORAZEPAM, LORAZEPAM
 LOSARTAN POTASSIUM AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 LOSARTAN POTASSIUM, LOSARTAN POTASSIUM
 LOXAPINE SUCCINATE, LOXAPINE SUCCINATE
 LUXIQ, BETAMETHASONE VALERATE
 MAPROTILINE HYDROCHLORIDE, MAPROTILINE HYDROCHLORIDE
 MECLOFENAMATE SODIUM, MECLOFENAMATE SODIUM
 MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
 MENTAX, BUTENAFINE HYDROCHLORIDE
 MERCAPTOPYRINE, MERCAPTOPYRINE
 MESALAMINE, MESALAMINE
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 METHIMAZOLE, METHIMAZOLE
 METHOTREXATE SODIUM, METHOTREXATE SODIUM
 METHYLDOPA AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 METHYLDOPA, METHYLDOPA
 METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
 METOLAZONE, METOLAZONE
 METOPROLOL TARTRATE, METOPROLOL TARTRATE
 MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE
 MIRTAZAPINE, MIRTAZAPINE
 MYCOPHENOLATE MOFETIL, MYCOPHENOLATE MOFETIL
 MYCOPHENOLIC ACID, MYCOPHENOLIC ACID
 NADOLOL, NADOLOL
 NEVIRAPINE, NEVIRAPINE
 NICARDIPINE HYDROCHLORIDE, NICARDIPINE HYDROCHLORIDE
 NIFEDIPINE, NIFEDIPINE
 NISOLDIPINE, NISOLDIPINE
 NITROFURANTOIN (MONOHYDRATE/MACROCRYSTALS), NITROFURANTOIN
 OLANZAPINE, OLANZAPINE
 OLMESARTAN MEDOXOMIL, OLMESARTAN MEDOXOMIL
 OLOPATADINE HYDROCHLORIDE, OLOPATADINE HYDROCHLORIDE
 OLUX E, CLOBETASOL PROPIONATE
 OLUX, CLOBETASOL PROPIONATE
 ONDANSETRON, ONDANSETRON

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** M ****

* MYLAN PHARMACEUTICALS INC
 PALIPERIDONE, PALIPERIDONE
 PAROXETINE HYDROCHLORIDE, PAROXETINE HYDROCHLORIDE
 PERPHENAZINE AND AMITRIPTYLINE HYDROCHLORIDE, AMITRIPTYLINE HYDROCHLORIDE
 PERPHENAZINE, PERPHENAZINE
 PHENYTEK, PHENYTOIN SODIUM
 POLYETHYLENE GLYCOL 3350, POLYETHYLENE GLYCOL 3350 (OTC)
 POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
 PRASUGREL, PRASUGREL HYDROCHLORIDE
 PRAZOSIN HYDROCHLORIDE, PRAZOSIN HYDROCHLORIDE
 PREDNISONE, PREDNISONE
 PROBENECID, PROBENECID
 PROCHLORPERAZINE MALEATE, PROCHLORPERAZINE MALEATE
 PROPAFENONE HYDROCHLORIDE, PROPAFENONE HYDROCHLORIDE
 PROPRANOLOL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 PROPRANOLOL HYDROCHLORIDE, PROPRANOLOL HYDROCHLORIDE
 QUINAPRIL HYDROCHLORIDE, QUINAPRIL HYDROCHLORIDE
 RASAGILINE MESYLATE, RASAGILINE MESYLATE
 RISPERIDONE, RISPERIDONE
 ROPINIROLE HYDROCHLORIDE, ROPINIROLE HYDROCHLORIDE
 SILDENAFIL CITRATE, SILDENAFIL CITRATE
 SOTALOL HYDROCHLORIDE, SOTALOL HYDROCHLORIDE
 SPIRONOLACTONE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 SPIRONOLACTONE, SPIRONOLACTONE
 SULINDAC, SULINDAC
 SUMATRIPTAN AND NAPROXEN SODIUM, NAPROXEN SODIUM
 SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
 SYMFI LO, EFAVIRENZ
 TACROLIMUS, TACROLIMUS
 TADALAFIL, TADALAFIL
 TAMOXIFEN CITRATE, TAMOXIFEN CITRATE
 TELMISARTAN AND AMLODIPINE, AMLODIPINE BESYLATE
 TELMISARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 TELMISARTAN, TELMISARTAN
 TEMAZEPAM, TEMAZEPAM
 TERAZOSIN HYDROCHLORIDE, TERAZOSIN HYDROCHLORIDE
 TERIFLUNOMIDE, TERIFLUNOMIDE
 TETRABENAZINE, TETRABENAZINE
 THIORIDAZINE HYDROCHLORIDE, THIORIDAZINE HYDROCHLORIDE
 THIOTHIXENE, THIOTHIXENE
 TIMOLOL MALEATE, TIMOLOL MALEATE
 TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE
 TOLMETIN SODIUM, TOLMETIN SODIUM
 TOLTERODINE TARTRATE, TOLTERODINE TARTRATE
 TRAMADOL HYDROCHLORIDE AND ACETAMINOPHEN, ACETAMINOPHEN
 TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE
 TRAVOPROST, TRAVOPROST
 TRETINOIN, TRETINOIN
 TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE
 TRIAMTERENE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 TRIFLUOPERAZINE HYDROCHLORIDE, TRIFLUOPERAZINE HYDROCHLORIDE
 TRILYTE, POLYETHYLENE GLYCOL 3350
 URSODIOL, URSODIOL
 VALSARTAN, VALSARTAN
 VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE
 VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE
 VUSION, MICONAZOLE NITRATE
 WIXELA INHUB, FLUTICASONE PROPIONATE
 ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE
 ZONALON, DOXEPIN HYDROCHLORIDE
 ZOVIRAX, ACYCLOVIR

MYLAN ASI

* MYLAN ASI LLC
 ACETAZOLAMIDE SODIUM, ACETAZOLAMIDE SODIUM
 ADENOSINE, ADENOSINE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** M ****

* MYLAN ASI LLC
 AZITHROMYCIN, AZITHROMYCIN
 GRANISETRON HYDROCHLORIDE, GRANISETRON HYDROCHLORIDE
 METOPROLOL TARTRATE, METOPROLOL TARTRATE
 POLYMYXIN B SULFATE, POLYMYXIN B SULFATE
 ROPIVACAINE HYDROCHLORIDE, ROPIVACAINE HYDROCHLORIDE
 SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE

MYLAN INSTITUTIONAL

* MYLAN INSTITUTIONAL INC
 MYLERAN, BUSULFAN
 SULFAMYLOL, MAFENIDE ACETATE

* MYLAN INSTITUTIONAL LLC
 ALOPRIM, ALLOPURINOL SODIUM
 AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE
 ARGATROBAN, ARGATROBAN
 AZACITIDINE, AZACITIDINE
 BIVALIRUDIN, BIVALIRUDIN
 CHLOROTHIAZIDE SODIUM, CHLOROTHIAZIDE SODIUM
 CIDOFOVIR, CIDOFOVIR
 COSYNTROPIN, COSYNTROPIN
 DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE
 DEXRAZOXANE HYDROCHLORIDE, DEXRAZOXANE HYDROCHLORIDE
 DIPHENHYDRAMINE HYDROCHLORIDE, DIPHENHYDRAMINE HYDROCHLORIDE
 DURACLON, CLONIDINE HYDROCHLORIDE
 ESMOLOL HYDROCHLORIDE, ESMOLOL HYDROCHLORIDE
 ETHACRYNATE SODIUM, ETHACRYNATE SODIUM
 FOMEPIZOLE, FOMEPIZOLE
 FULVESTRANT, FULVESTRANT
 HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
 IBUTILIDE FUMARATE, IBUTILIDE FUMARATE
 ISOSULFAN BLUE, ISOSULFAN BLUE
 KETAMINE HYDROCHLORIDE, KETAMINE HYDROCHLORIDE
 MEFOXIN IN PLASTIC CONTAINER, CEFOXITIN SODIUM
 MELPHALAN HYDROCHLORIDE, MELPHALAN HYDROCHLORIDE
 METHADONE HYDROCHLORIDE, METHADONE HYDROCHLORIDE
 METHOCARBAMOL, METHOCARBAMOL
 NALOXONE HYDROCHLORIDE, NALOXONE HYDROCHLORIDE
 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

* MYLAN LABORATORIES LTD
 NEVIRAPINE, NEVIRAPINE

MYLAN LABS LTD

* MYLAN LABORATORIES LTD
 ADENOSINE, ADENOSINE
 AMIFOSTINE, AMIFOSTINE
 AMPICILLIN AND SULBACTAM, AMPICILLIN SODIUM
 AMPICILLIN SODIUM, AMPICILLIN SODIUM
 AZITHROMYCIN, AZITHROMYCIN
 BACLOFEN, BACLOFEN
 CAPREOMYCIN SULFATE, CAPREOMYCIN SULFATE
 CASPOFUNGIN ACETATE, CASPOFUNGIN ACETATE
 CIMDUO, LAMIVUDINE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** M ****

* MYLAN LABORATORIES LTD
 CLADRIBINE, CLADRIBINE
 CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
 CLOFARABINE, CLOFARABINE
 CYANOCOBALAMIN, CYANOCOBALAMIN
 CYTARABINE, CYTARABINE
 DACTINOMYCIN, DACTINOMYCIN
 DAPTOMYCIN, DAPTOMYCIN
 DESOGESTREL AND ETHINYL ESTRADIOL, DESOGESTREL
 DEXAMETHASONE SODIUM PHOSPHATE, DEXAMETHASONE SODIUM PHOSPHATE
 DOCETAXEL, DOCETAXEL
 DOXORUBICIN HYDROCHLORIDE, DOXORUBICIN HYDROCHLORIDE
 DOXYCYCLINE, DOXYCYCLINE HYCLATE
 DROSPIRENONE AND ETHINYL ESTRADIOL, DROSPIRENONE
 EPTIFIBATIDE, EPTIFIBATIDE
 ESMOLOL HYDROCHLORIDE, ESMOLOL HYDROCHLORIDE
 ESOMEPRAZOLE SODIUM, ESOMEPRAZOLE SODIUM
 ESTRADIOL AND NORETHINDRONE ACETATE, ESTRADIOL
 ETHYNODIOL DIACETATE AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
 ETOMIDATE, ETOMIDATE
 ETOPOSIDE, ETOPOSIDE
 FAMOTIDINE PRESERVATIVE FREE, FAMOTIDINE
 FAMOTIDINE, FAMOTIDINE
 FLUDARABINE PHOSPHATE, FLUDARABINE PHOSPHATE
 FLUMAZENIL, FLUMAZENIL
 FLUOROURACIL, FLUOROURACIL
 FLUPHENAZINE DECANOATE, FLUPHENAZINE DECANOATE
 FOSAPREPITANT DIMEGLUMINE, FOSAPREPITANT DIMEGLUMINE
 FOSPHENYTOIN SODIUM, FOSPHENYTOIN SODIUM
 GANCICLOVIR SODIUM, GANCICLOVIR SODIUM
 GEMCITABINE HYDROCHLORIDE, GEMCITABINE HYDROCHLORIDE
 GRANISETRON HYDROCHLORIDE, GRANISETRON HYDROCHLORIDE
 HALOPERIDOL DECANOATE, HALOPERIDOL DECANOATE
 HALOPERIDOL, HALOPERIDOL LACTATE
 HEPARIN SODIUM, HEPARIN SODIUM
 IBANDRONATE SODIUM, IBANDRONATE SODIUM
 LAMIVUDINE, LAMIVUDINE
 LEUCOVORIN CALCIUM PRESERVATIVE FREE, LEUCOVORIN CALCIUM
 LEVETIRACETAM, LEVETIRACETAM
 LEVOFLOXACIN, LEVOFLOXACIN
 LEVONORGESTREL AND ETHINYL ESTRADIOL AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
 LEVONORGESTREL AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
 LEVONORGESTREL, LEVONORGESTREL
 LEVONORGESTREL, LEVONORGESTREL (OTC)
 LINEZOLID, LINEZOLID
 MAGNESIUM SULFATE IN DEXTROSE 5% IN PLASTIC CONTAINER, MAGNESIUM SULFATE
 MAGNESIUM SULFATE IN PLASTIC CONTAINER, MAGNESIUM SULFATE
 MEDROXYPROGESTERONE ACETATE, MEDROXYPROGESTERONE ACETATE
 METHOTREXATE SODIUM PRESERVATIVE FREE, METHOTREXATE SODIUM
 METRONIDAZOLE IN PLASTIC CONTAINER, METRONIDAZOLE
 MITOMYCIN, MITOMYCIN
 MOXIFLOXACIN HYDROCHLORIDE IN SODIUM CHLORIDE 0.8% IN PLASTIC CONTAINER, MOXIFLOXACIN
 MYCOPHENOLATE MOFETIL HYDROCHLORIDE, MYCOPHENOLATE MOFETIL HYDROCHLORIDE
 NOREPINEPHRINE BITARTRATE, NOREPINEPHRINE BITARTRATE
 NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL AND FERROUS FUMARATE, ETHINYL ESTRADIOL
 NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
 NORETHINDRONE ACETATE, NORETHINDRONE ACETATE
 NORETHINDRONE AND ETHINYL ESTRADIOL AND FERROUS FUMARATE, ETHINYL ESTRADIOL
 NORETHINDRONE, NORETHINDRONE
 NORGESTIMATE AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
 ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
 OXALIPLATIN, OXALIPLATIN
 PACLITAXEL, PACLITAXEL
 PAMIDRONATE DISODIUM, PAMIDRONATE DISODIUM
 PARICALCITOL, PARICALCITOL

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** M ****

* MYLAN LABORATORIES LTD

PIPERACILLIN AND TAZOBACTAM, PIPERACILLIN SODIUM
 PROCHLORPERAZINE EDISYLATE, PROCHLORPERAZINE EDISYLATE
 RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
 RIFAMPIN, RIFAMPIN
 SODIUM NITROPRUSSIDE, SODIUM NITROPRUSSIDE
 SULFAMETHOXAZOLE AND TRIMETHOPRIM, SULFAMETHOXAZOLE
 SYMFI, EFAVIRENZ
 TOBRAMYCIN SULFATE, TOBRAMYCIN SULFATE
 TOPOTECAN HYDROCHLORIDE, TOPOTECAN HYDROCHLORIDE
 VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE
 VECURONIUM BROMIDE, VECURONIUM BROMIDE
 ZOLEDRONIC ACID, ZOLEDRONIC ACID

MYLAN PHARMS INC

* MYLAN PHARMACEUTICALS INC

ABACAVIR SULFATE, ABACAVIR SULFATE
 ACYCLOVIR, ACYCLOVIR
 AMNESTEEM, ISOTRETINOIN
 ARMODAFINIL, ARMODAFINIL
 ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM
 AVITA, TRETINOIN
 BACLOFEN, BACLOFEN
 CHLORDIAZEPOXIDE AND AMITRIPTYLINE HYDROCHLORIDE, AMITRIPTYLINE HYDROCHLORIDE
 CLINDAMYCIN PHOSPHATE AND BENZOYL PEROXIDE, BENZOYL PEROXIDE
 CYCLOBENZAPRINE HYDROCHLORIDE, CYCLOBENZAPRINE HYDROCHLORIDE
 DICLOFENAC SODIUM, DICLOFENAC SODIUM
 DIGOXIN, DIGOXIN
 DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
 DOXEPIN HYDROCHLORIDE, DOXEPIN HYDROCHLORIDE
 EPROSARTAN MESYLATE, EPROSARTAN MESYLATE
 ESZOPICLONE, ESZOPICLONE
 FENOFIBRATE (MICRONIZED), FENOFIBRATE
 FENOFIBRATE, FENOFIBRATE
 FLURAZEPAM HYDROCHLORIDE, FLURAZEPAM HYDROCHLORIDE
 GABAPENTIN, GABAPENTIN
 ITRACONAZOLE, ITRACONAZOLE
 LANSOPRAZOLE, LANSOPRAZOLE
 LITHIUM CARBONATE, LITHIUM CARBONATE
 MAXZIDE, HYDROCHLOROTHIAZIDE
 MAXZIDE-25, HYDROCHLOROTHIAZIDE
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 METOPROLOL SUCCINATE, METOPROLOL SUCCINATE
 MIDODRINE HYDROCHLORIDE, MIDODRINE HYDROCHLORIDE
 MODAFINIL, MODAFINIL
 NABUMETONE, NABUMETONE
 NEVIRAPINE, NEVIRAPINE
 PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
 PHENYTOIN, PHENYTOIN
 PINDOLOL, PINDOLOL
 PRAVASTATIN SODIUM, PRAVASTATIN SODIUM
 PREDNISOLONE SODIUM PHOSPHATE, PREDNISOLONE SODIUM PHOSPHATE
 RILUZOLE, RILUZOLE
 RIZATRIPTAN BENZOATE, RIZATRIPTAN BENZOATE
 SILDENAFIL CITRATE, SILDENAFIL CITRATE
 TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE
 TOLAZAMIDE, TOLAZAMIDE
 TOLBUTAMIDE, TOLBUTAMIDE
 VALACYCLOVIR HYDROCHLORIDE, VALACYCLOVIR HYDROCHLORIDE
 VALSARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 VORICONAZOLE, VORICONAZOLE
 ZIDOVUDINE, ZIDOVUDINE

* MYLAN PHARMACEUTICALS INC.

FLUVASTATIN SODIUM, FLUVASTATIN SODIUM
 NIZATIDINE, NIZATIDINE

MYLAN SPECIALITY LP

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** M ****

* MYLAN SPECIALTY LP
 ACCUNEB, ALBUTEROL SULFATE
 ANADROL-50, OXYMETHOLONE
 ASTELIN, AZELASTINE HYDROCHLORIDE
 ASTEPRO, AZELASTINE HYDROCHLORIDE
 AVC, SULFANILAMIDE
 BUTISOL SODIUM, BUTABARBITAL SODIUM
 CESAMET, NABILONE
 COLYTE WITH FLAVOR PACKS, POLYETHYLENE GLYCOL 3350
 CORTIFOAM, HYDROCORTISONE ACETATE
 DEMADEX, TORSEMIDE
 DEPEN, PENICILLAMINE
 DIPENTUM, OLSALAZINE SODIUM
 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
 MUSE, ALPROSTADIL
 PROCTOFOAM HC, HYDROCORTISONE ACETATE
 ROWASA, MESALAMINE
 SFROWASA, MESALAMINE
 SOMA, CARISOPRODOL
 TOBI PODHALER, TOBRAMYCIN
 TOBI, TOBRAMYCIN

MYLAN SPECLT

* MYLAN SPECIALTY LP
 PERFORMIST, FORMOTEROL FUMARATE

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
 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
 LEVONORGESTREL, LEVONORGESTREL (OTC)
 NORGESTIMATE AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL

NABRIVA

* NABRIVA THERAPEUTICS IRELAND DAC
 XENLETA, LEFAMULIN ACETATE

NALPROPION

* NALPROPION PHARMACEUTICALS LLC
 CONTRAVE, BUPROPION HYDROCHLORIDE

NANG KUANG PHARM CO

* NANG KUANG PHARMACEUTICAL CO LTD
 LINEZOLID, LINEZOLID

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** N ******NANJING KING-FRIEND**

- * NANJING KING-FRIEND BIOCHEMICAL PHARMACEUTICAL CO LTD
ENOXAPARIN SODIUM (PRESERVATIVE FREE), ENOXAPARIN SODIUM
HEPARIN SODIUM, HEPARIN SODIUM

NAPO PHARMS INC

- * NAPO PHARMACEUTICALS INC
MYTESI, CROFELEMER

NATCO PHARMA

- * NATCO PHARMA LTD
GRANISETRON HYDROCHLORIDE, GRANISETRON HYDROCHLORIDE

NATCO PHARMA LTD

- * NATCO PHARMA LIMITED
CHLOROQUINE PHOSPHATE, CHLOROQUINE PHOSPHATE
- * NATCO PHARMA LTD
ALPRAZOLAM, ALPRAZOLAM
ANASTROZOLE, ANASTROZOLE
ARMODAFINIL, ARMODAFINIL
AZACITIDINE, AZACITIDINE
BOSENTAN, BOSENTAN
CARISOPRODOL, CARISOPRODOL
CHLOROQUINE PHOSPHATE, CHLOROQUINE PHOSPHATE
ERLOTINIB HYDROCHLORIDE, ERLOTINIB HYDROCHLORIDE
GLYCOPYRROLATE, GLYCOPYRROLATE
IMATINIB MESYLATE, IMATINIB MESYLATE
LANSOPRAZOLE, LANSOPRAZOLE
LANSOPRAZOLE, LANSOPRAZOLE (OTC)
LANTHANUM CARBONATE, LANTHANUM CARBONATE
LETROZOLE, LETROZOLE
ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
OSELTAMIVIR PHOSPHATE, OSELTAMIVIR PHOSPHATE
RIZATRIPTAN BENZOATE, RIZATRIPTAN BENZOATE
TRIHENXYPHENIDYL HYDROCHLORIDE, TRIHENXYPHENIDYL HYDROCHLORIDE

NAVINTA LLC

- * NAVINTA LLC
BENZTROPINE MESYLATE, BENZTROPINE MESYLATE
CARMUSTINE, CARMUSTINE
FAMOTIDINE, FAMOTIDINE
FOMEPIZOLE, FOMEPIZOLE
HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
INDOMETHACIN SODIUM, INDOMETHACIN SODIUM
METHOCARBAMOL, METHOCARBAMOL
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

NEOPHARMA

- * NEOPHARMA INC
ALENDRONATE SODIUM, ALENDRONATE SODIUM
AMOXIL, AMOXICILLIN
ANASTROZOLE, ANASTROZOLE
AUGMENTIN '125', AMOXICILLIN
AUGMENTIN '250', AMOXICILLIN
AUGMENTIN '875', AMOXICILLIN
AUGMENTIN XR, AMOXICILLIN
IRBESARTAN, IRBESARTAN
IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE
LAROTID, AMOXICILLIN
LEVOCETIRIZINE DIHYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE
PIOGLITAZONE HYDROCHLORIDE, PIOGLITAZONE HYDROCHLORIDE

NEOS THERAP INC

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** N ****

* NEOS THERAPEUTICS INC
HYDROCODONE POLISTIREX AND CHLORPHENIRAMNE POLISTIREX, CHLORPHENIRAMINE POLISTIREX

NEOS THERAPS

* NEOS THERAPEUTICS
ADZENYS XR-ODT, AMPHETAMINE

NEOS THERAPS INC

* NEOS THERAPEUTICS INC
ADZENYS ER, AMPHETAMINE
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
BUDESONIDE, BUDESONIDE

NESHER PHARMS

* NESHER PHARMACEUTICALS USA LLC
DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE
DEXTROAMPHETAMINE SULFATE, DEXTROAMPHETAMINE SULFATE
ISOSORBIDE MONONITRATE, ISOSORBIDE MONONITRATE
MICRO-K 10, POTASSIUM CHLORIDE
MICRO-K, POTASSIUM CHLORIDE
MORPHINE SULFATE, MORPHINE SULFATE
NYSTATIN, NYSTATIN
OSELTAMIVIR PHOSPHATE, OSELTAMIVIR PHOSPHATE
OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN
OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE

NEURELIS INC

* NEURELIS INC
VALTOCO, DIAZEPAM

NEUROCRINE

* NEUROCRINE BIOSCIENCES INC
INGREZZA, VALBENZAZINE TOSYLATE

NEXGEN PHARMA

* NEXGEN PHARMA INC
BUTALBITAL AND ACETAMINOPHEN, ACETAMINOPHEN
BUTALBITAL, ACETAMINOPHEN AND CAFFEINE, ACETAMINOPHEN
CHENODIOL, CHENODIOL
GLYCOPYRROLATE, GLYCOPYRROLATE
MECAMYLAMINE HYDROCHLORIDE, MECAMYLAMINE HYDROCHLORIDE
POLYETHYLENE GLYCOL 3350, POLYETHYLENE GLYCOL 3350 (OTC)

NEXGEN PHARMA INC

* NEXGEN PHARMA INC
BUTALBITAL, ACETAMINOPHEN AND CAFFEINE, ACETAMINOPHEN
BUTALBITAL, ACETAMINOPHEN, CAFFEINE AND CODEINE PHOSPHATE, ACETAMINOPHEN
BUTALBITAL, ASPIRIN, CAFFEINE, AND CODEINE PHOSPHATE, ASPIRIN
NABUMETONE, NABUMETONE

NEXTWAVE

* NEXTWAVE PHARMACEUTICALS INC A SUB OF TRIS PHARMA INC
QUILLIVANT XR, METHYLPHENIDATE HYDROCHLORIDE

NEXTWAVE PHARMS

* NEXTWAVE PHARMACEUTICALS INC
QUILLICHEW ER, METHYLPHENIDATE HYDROCHLORIDE

NEXUS PHARMS

* NEXUS PHARMACEUTICALS INC
ARSENIC TRIOXIDE, ARSENIC TRIOXIDE
BUSULFAN, BUSULFAN
COLISTIMETHATE SODIUM, COLISTIMETHATE SODIUM
DICYCLOMINE HYDROCHLORIDE, DICYCLOMINE HYDROCHLORIDE
ISOPROTERENOL HYDROCHLORIDE, ISOPROTERENOL HYDROCHLORIDE
PROCAINAMIDE HYDROCHLORIDE, PROCAINAMIDE HYDROCHLORIDE
PROCHLORPERAZINE EDISYLATE, PROCHLORPERAZINE EDISYLATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** N ****

* NEXUS PHARMACEUTICALS INC
SODIUM NITROPRUSSIDE, SODIUM NITROPRUSSIDE
TESTOSTERONE ENANTHATE, TESTOSTERONE ENANTHATE

NIAGARA PHARMS

* NIAGARA PHARMACEUTICALS INC
PUR-WASH, PURIFIED WATER (OTC)

NODEN PHARMA

* NODEN PHARMA DAC
TEKTURNA HCT, ALISKIREN HEMIFUMARATE
TEKTURNA, ALISKIREN HEMIFUMARATE

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

NORTHSTAR MEDICAL

* NORTHSTAR MEDICAL RADIOISOTOPES LLC
RADIOGENIX SYSTEM, TECHNETIUM TC-99M SODIUM PERTECHNETATE GENERATOR

NORTON WATERFORD

* NORTON WATERFORD LTD
QVAR REDIHALER, BECLOMETHASONE DIPROPIONATE

NOSTRUM LABS INC

* NOSTRUM LABORATORIES INC
ACETAZOLAMIDE, ACETAZOLAMIDE
CALCIUM ACETATE, CALCIUM ACETATE
CARISOPRODOL, CARISOPRODOL
CLARITHROMYCIN, CLARITHROMYCIN
DAPSONE, DAPSONE
ELIXOPHYLLIN, THEOPHYLLINE
ENALAPRIL MALEATE AND HYDROCHLOROTHIAZIDE, ENALAPRIL MALEATE
ENALAPRIL MALEATE, ENALAPRIL MALEATE
HYDROMORPHONE HYDROCHLORIDE, HYDROMORPHONE HYDROCHLORIDE
METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
MORPHINE SULFATE, MORPHINE SULFATE
NABUMETONE, NABUMETONE
NITROFURANTOIN, NITROFURANTOIN
OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN
PINDOLOL, PINDOLOL
PIROXICAM, PIROXICAM
PROMETHAZINE HYDROCHLORIDE AND CODEINE PHOSPHATE, CODEINE PHOSPHATE
PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
THEOPHYLLINE, THEOPHYLLINE
VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE

NOSTRUM PHARMS LLC

* NOSTRUM PHARMACEUTICALS LLC
METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
THEOCHRON, THEOPHYLLINE

NOVA LABS LTD

* NOVA LABORATORIES LTD
PURIXAN, MERCAPTOPYRINE

NOVADAQ TECH

* NOVADAQ TECHNOLOGIES ULC
SPY AGENT GREEN KIT, INDOCYANINE GREEN

NOVARTIS

* NOVARTIS PHARMACEUTICALS CORP

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** N **

* NOVARTIS PHARMACEUTICALS CORP
 AFINITOR, EVEROLIMUS
 ALOMIDE, LODOXAMIDE TROMETHAMINE
 ARGATROBAN, ARGATROBAN
 ARRANON, NELARABINE
 AZOPT, BRINZOLAMIDE
 BETOPTIC S, BETAXOLOL HYDROCHLORIDE
 CILOXAN, CIPROFLOXACIN HYDROCHLORIDE
 CIPRO HC, CIPROFLOXACIN HYDROCHLORIDE
 CIPRODEX, CIPROFLOXACIN
 COARTEM, ARTEMETHER
 DESFERAL, DEFEROXAMINE MESYLATE
 DIOVAN HCT, HYDROCHLOROTHIAZIDE
 DIOVAN, VALSARTAN
 DUREZOL, DIFLUPREDNATE
 EGATEN, TRICLABENDAZOLE
 EXELON, RIVASTIGMINE
 EXFORGE HCT, AMLODIPINE BESYLATE
 EXFORGE, AMLODIPINE BESYLATE
 EXJADE, DEFERASIROX
 FOCALIN XR, DEXMETHYLPHENIDATE HYDROCHLORIDE
 FOCALIN, DEXMETHYLPHENIDATE HYDROCHLORIDE
 GILENYA, FINGOLIMOD HYDROCHLORIDE
 GLEEVEC, IMATINIB MESYLATE
 HYCAMTIN, TOPOTECAN HYDROCHLORIDE
 ILEVRO, NEPAFENAC
 IOPIDINE, APRACLOLONIDINE HYDROCHLORIDE
 ISOPTO CARPINE, PILOCARPINE HYDROCHLORIDE
 JADENU SPRINKLE, DEFERASIROX
 KISQALI FEMARA CO-PACK (COPACKAGED), LETROZOLE
 KISQALI, RIBOCICLIB SUCCINATE
 LAMISIL, TERBINAFINE HYDROCHLORIDE
 LESCOL XL, FLUVASTATIN SODIUM
 LOTREL, AMLODIPINE BESYLATE
 MAXIDEX, DEXAMETHASONE
 MAXITROL, DEXAMETHASONE
 MAYZENT, SIPONIMOD FUMARIC ACID
 MEKINIST, TRAMETINIB DIMETHYL SULFOXIDE
 MOXEZA, MOXIFLOXACIN HYDROCHLORIDE
 MYDRIACYL, TROPICAMIDE
 MYFORTIC, MYCOPHENOLIC ACID
 NATACYN, NATAMYCIN
 NEORAL, CYCLOSPORINE
 NEVANAC, NEPAFENAC
 OMNIPRED, PREDNISOLONE ACETATE
 PATADAY, OLOPATADINE HYDROCHLORIDE
 PATANASE, OLOPATADINE HYDROCHLORIDE
 PATANOL, OLOPATADINE HYDROCHLORIDE
 PAZEO, OLOPATADINE HYDROCHLORIDE
 PIQRAY, ALPELISIB
 PROMACTA KIT, ELTROMBOPAG OLAMINE
 PROMACTA, ELTROMBOPAG OLAMINE
 RECLAST, ZOLEDRONIC ACID
 RITALIN LA, METHYLPHENIDATE HYDROCHLORIDE
 RITALIN, METHYLPHENIDATE HYDROCHLORIDE
 RYDAPT, MIDOSTAURIN
 SANDIMMUNE, CYCLOSPORINE
 SANDOSTATIN LAR, OCTREOTIDE ACETATE
 SANDOSTATIN, OCTREOTIDE ACETATE
 SIMBRINZA, BRIMONIDINE TARTRATE
 TAFINLAR, DABRAFENIB MESYLATE
 TASIGNA, NILOTINIB HYDROCHLORIDE
 TEGRETOL, CARBAMAZEPINE
 TEGRETOL-XR, CARBAMAZEPINE
 TOBRADEX, DEXAMETHASONE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** N ****

* NOVARTIS PHARMACEUTICALS CORP
 TOBEX, TOBRAMYCIN
 TRAVATAN Z, TRAVOPROST
 TRIESENCE, TRIAMCINOLONE ACETONIDE
 TRILEPTAL, OXCARBAZEPINE
 TYKERB, LAPATINIB DITOSYLATE
 VIGAMOX, MOXIFLOXACIN HYDROCHLORIDE
 VIVELLE-DOT, ESTRADIOL
 VOLTAREN, DICLOFENAC SODIUM
 VOTRIENT, PAZOPANIB HYDROCHLORIDE
 XIIDRA, LIFITEGRAST
 ZOFRAN ODT, ONDANSETRON
 ZOFRAN, ONDANSETRON HYDROCHLORIDE
 ZOMETA, ZOLEDRONIC ACID
 ZORTRESS, EVEROLIMUS
 ZYKADIA, CERITINIB

NOVARTIS PHARM

* NOVARTIS PHARMACEUTICAL CORP
 AFINITOR DISPERZ, EVEROLIMUS

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 LTD
 ACETAZOLAMIDE, ACETAZOLAMIDE
 CARISOPRODOL AND ASPIRIN, ASPIRIN
 CARISOPRODOL, CARISOPRODOL
 CICLOPIROX, CICLOPIROX
 CLOBETASOL PROPIONATE (EMOLLIENT), CLOBETASOL PROPIONATE
 CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
 CYPROHEPTADINE HYDROCHLORIDE, CYPROHEPTADINE HYDROCHLORIDE
 DESIPRAMINE HYDROCHLORIDE, DESIPRAMINE HYDROCHLORIDE
 DESMOPRESSIN ACETATE, DESMOPRESSIN ACETATE
 DESOGESTREL AND ETHINYL ESTRADIOL, DESOGESTREL
 DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
 ESTRADIOL AND NORETHINDRONE ACETATE, ESTRADIOL
 HER STYLE, LEVONORGESTREL (OTC)
 INDOMETHACIN, INDOMETHACIN
 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
 LIDOCAINE, LIDOCAINE
 LO-MALMOREDE, ETHINYL ESTRADIOL
 MAFENIDE ACETATE, MAFENIDE ACETATE
 MALMOREDE, ETHINYL ESTRADIOL
 MELAMISA, DROSPIRENONE
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 METOPROLOL SUCCINATE, METOPROLOL SUCCINATE
 NADOLOL, NADOLOL
 NIFEDIPINE, NIFEDIPINE
 NORETHINDRONE ACETATE, NORETHINDRONE ACETATE
 NORETHINDRONE, NORETHINDRONE
 OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE
 PIMTREA, DESOGESTREL
 PRIMAQUINE PHOSPHATE, PRIMAQUINE PHOSPHATE
 PROBENECID AND COLCHICINE, COLCHICINE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** N ****

* NOVAST LABORATORIES LTD
 QUETIAPINE FUMARATE, QUETIAPINE FUMARATE
 QUININE SULFATE, QUININE SULFATE
 SETLAKIN, ETHINYL ESTRADIOL
 TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE
 TOLCAPONE, TOLCAPONE
 TOPOTECAN HYDROCHLORIDE, TOPOTECAN HYDROCHLORIDE
 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
 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
 CALCIPOTRIENE, CALCIPOTRIENE
 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
 DICLOFENAC SODIUM, DICLOFENAC SODIUM
 DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
 FAMOTIDINE, FAMOTIDINE
 FLUCYTOSINE, FLUCYTOSINE
 FLUOCINONIDE, FLUOCINONIDE
 HOMATROPINE METHYLBROMIDE AND HYDROCODONE BITARTRATE, HOMATROPINE METHYLBROMIDE
 HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
 HYDROCODONE BITARTRATE AND HOMATROPINE METHYLBROMIDE, HOMATROPINE METHYLBROMIDE
 LEVONORGESTREL, LEVONORGESTREL (OTC)
 LINEZOLID, LINEZOLID
 METHYLERGONOVINE MALEATE, METHYLERGONOVINE MALEATE
 METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
 METOCLOPRAMIDE HYDROCHLORIDE, METOCLOPRAMIDE HYDROCHLORIDE
 MISOPROSTOL, MISOPROSTOL
 MORPHINE SULFATE, MORPHINE SULFATE
 MOXIFLOXACIN HYDROCHLORIDE, MOXIFLOXACIN HYDROCHLORIDE
 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
 TINIDAZOLE, TINIDAZOLE
 TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE
 TRIMETHOPRIM, TRIMETHOPRIM
 VORICONAZOLE, VORICONAZOLE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** N ****

* NOVEL LABORATORIES INC
ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE

NOVELGENIX THERAPS

* NOVELGENIX THERAPEUTICS PVT LTD
METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
NAPROXEN SODIUM, NAPROXEN SODIUM (OTC)
POLYETHYLENE GLYCOL 3350, POLYETHYLENE GLYCOL 3350 (OTC)
SODIUM POLYSTYRENE SULFONATE, SODIUM POLYSTYRENE SULFONATE
VALPROIC ACID, VALPROIC ACID

NOVEN

* NOVEN PHARMACEUTICALS INC
MINIVELLE, ESTRADIOL

NOVEN PHARMS INC

* NOVEN PHARMACEUTICALS INC
COMBIPATCH, ESTRADIOL
DAYTRANA, METHYLPHENIDATE

NOVITIUM PHARMA

* NOVITIUM PHARMA LLC
ACETAZOLAMIDE, ACETAZOLAMIDE
CHLORZOXAZONE, CHLORZOXAZONE
CLOTRIMAZOLE, CLOTRIMAZOLE
DAPSONE, DAPSONE
LEVOCARNITINE SF, LEVOCARNITINE
LEVOCARNITINE, LEVOCARNITINE
NITISINONE, NITISINONE
OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE
PRAZOSIN HYDROCHLORIDE, PRAZOSIN HYDROCHLORIDE
PREDNISONE, PREDNISONE
PYRIDOSTIGMINE BROMIDE, PYRIDOSTIGMINE BROMIDE
RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
SILDENAFIL CITRATE, SILDENAFIL CITRATE
SIROLIMUS, SIROLIMUS
THIOTHIXENE, THIOTHIXENE
TRIHEXYPHENIDYL HYDROCHLORIDE, TRIHEXYPHENIDYL HYDROCHLORIDE

NOVO

* NOVO NORDISK INC
FIASP FLEXTOUCH, INSULIN ASPART
FIASP PENFILL, INSULIN ASPART
FIASP, INSULIN ASPART
MACRILEN, MACIMORELIN ACETATE
OZEMPIC, SEMAGLUTIDE
RYBELSUS, SEMAGLUTIDE
SAXENDA, LIRAGLUTIDE RECOMBINANT
TRESIBA, INSULIN DEGLUDEC
XULTOPHY 100/3.6, INSULIN DEGLUDEC

NOVO NORDISK

* NOVO NORDISK PHARMACEUTICALS INC
GLUCAGEN, GLUCAGON HYDROCHLORIDE

NOVO NORDISK INC

* NOVO NORDISK INC
LEVEMIR FLEXTOUCH, INSULIN DETEMIR RECOMBINANT
LEVEMIR, INSULIN DETEMIR RECOMBINANT
NORDITROPIN FLEXPRO, SOMATROPIN
NOVOLIN 70/30, INSULIN RECOMBINANT HUMAN (OTC)
NOVOLIN N, INSULIN SUSP ISOPHANE RECOMBINANT HUMAN (OTC)
NOVOLIN R, INSULIN RECOMBINANT HUMAN (OTC)
NOVOLOG FLEXPEN, INSULIN ASPART RECOMBINANT
NOVOLOG MIX 70/30 FLEXPEN, INSULIN ASPART PROTAMINE RECOMBINANT
NOVOLOG MIX 70/30, INSULIN ASPART PROTAMINE RECOMBINANT
NOVOLOG PENFILL, INSULIN ASPART RECOMBINANT
NOVOLOG, INSULIN ASPART RECOMBINANT
VAGIFEM, ESTRADIOL
VICTOZA, LIRAGLUTIDE RECOMBINANT

NOVOCOL INC

* NOVOCOL INC

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** N ****

* NOVOCOL INC
DYCLOPRO, DYCLONINE HYDROCHLORIDE

NPS PHARMS INC

* NPS PHARMACEUTICALS INC
GATEX KIT, TEDUGLUTIDE RECOMBINANT

NUVO PHARM

* NUVO PHARMACEUTICAL INC
DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE
DEXTROAMPHETAMINE SULFATE, DEXTROAMPHETAMINE SULFATE
HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE
OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
PHENTERMINE HYDROCHLORIDE, PHENTERMINE HYDROCHLORIDE

NUVO PHARMS INC

* NUVO PHAMACEUTICALS INC
BUTALBITAL, ACETAMINOPHEN AND CAFFEINE, ACETAMINOPHEN
FOLIC ACID, FOLIC ACID
HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE
POLYETHYLENE GLYCOL 3350, POLYETHYLENE GLYCOL 3350 (OTC)
SODIUM POLYSTYRENE SULFONATE, SODIUM POLYSTYRENE SULFONATE

NXDC

* NX DEVELOPMENT CORP
GLEOLAN, AMINOLEVULINIC ACID HYDROCHLORIDE

**** O ******OAK PHARMS**

* OAK PHARMACEUTICALS INC
NEMBUTAL SODIUM, PENTOBARBITAL SODIUM
XYLOCAINE, LIDOCAINE HYDROCHLORIDE

OAK PHARMS AKORN

* OAK PHARMACEUTICALS INC SUB AKORN INC
COGENTIN, BENZTROPINE MESYLATE
DIURIL, CHLOROTHIAZIDE SODIUM

OAK PHARMS INC

* OAK PHARMACEUTICALS INC
ZIOPTAN, TAFLUPROST
* OAK PHARMACEUTICALS INC SUBSIDIARY OF AKORN INC
AZASITE, AZITHROMYCIN
BETIMOL, TIMOLOL
COSOPT PF, DORZOLAMIDE HYDROCHLORIDE
COSOPT, DORZOLAMIDE HYDROCHLORIDE
XOPENEX, LEVALBUTEROL HYDROCHLORIDE

OAKLOCK LLC

* OAKLOCK LLC
DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE

OCULAR THERAPEUTIX

* OCULAR THERAPEUTIX INC
DEXTENZA, DEXAMETHASONE

ODYSSEY PHARMS

* ODYSSEY PHARMACEUTICALS INC
ANTABUSE, DISULFIRAM
URECHOLINE, BETHANECHOL CHLORIDE

OHM

* OHM CORP
IBUPROFEN, IBUPROFEN (OTC)

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
VALSARTAN, VALSARTAN

OLTA PHARMS

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** O **

* OLTA PHARMACEUTICALS CORP
LINDANE, LINDANE

OMEROS

* OMEROS CORP
OMIDRIA, KETOROLAC TROMETHAMINE

ONY

* ONY INC
INFASURF PRESERVATIVE FREE, CALFACTANT

ONYX THERAP

* ONYX THERAPEUTICS INC A WHOLLY OWNED SUB OF AMGEN INC
KYPROLIS, CARFILZOMIB

OPKO IRELAND GLOBAL

* OPKO IRELAND GLOBAL HOLDINGS LTD
RAYALDEE, CALCIFEDIOL

OPTINOSE US INC

* OPTINOSE US INC
XHANCE, FLUTICASONE PROPIONATE

ORAPHARMA

* ORAPHARMA INC
ARESTIN, MINOCYCLINE HYDROCHLORIDE

ORCHID HLTHCARE

* ORCHID HEALTHCARE
AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
ARIPIPIRAZOLE, ARIPIPIRAZOLE
CEFADROXIL, CEFADROXIL/CEFADROXIL HEMIHYDRATE
CEFDINIR, CEFDINIR
CEFPODOXIME PROXETIL, CEFPODOXIME PROXETIL
CEFPROZIL, CEFPROZIL
CEFUROXIME AXETIL, CEFUROXIME AXETIL
CEPHALEXIN, CEPHALEXIN
CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
CETIRIZINE HYDROCHLORIDE HIVES, CETIRIZINE HYDROCHLORIDE (OTC)
DESLORATADINE, DESLORATADINE
DIVALPROEX SODIUM, DIVALPROEX SODIUM
ESZOPICLONE, ESZOPICLONE
FELODIPINE, FELODIPINE
GEMIFLOXACIN MESYLATE, GEMIFLOXACIN MESYLATE
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 SUB MERCK

* ORGANON USA INC A SUB OF MERCK AND CO INC
BRIDION, SUGAMMADEX SODIUM
NUVARING, ETHINYL ESTRADIOL

ORGANON USA INC

* ORGANON USA INC
FOLLISTIM AQ, FOLLITROPIN ALFA/BETA

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** O **

* ORGANON USA INC

GANIRELIX ACETATE, GANIRELIX ACETATE
 NEXPLANON, ETNOGESTREL
 PREGNYL, GONADOTROPIN, CHORIONIC
 REMERON SOLTAB, MIRTAZAPINE
 REMERON, MIRTAZAPINE

ORIENT PHARMA CO LTD

* ORIENT PHARMA CO LTD

CARISOPRODOL, CARISOPRODOL
 MIGLITOL, MIGLITOL
 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

ORIT LABS LLC

* ORIT LABORATORIES LLC

BENZONATATE, BENZONATATE
 BISOPROLOL FUMARATE, BISOPROLOL FUMARATE
 CLINDAMYCIN PALMITATE HYDROCHLORIDE, CLINDAMYCIN PALMITATE HYDROCHLORIDE
 CYCLOBENZAPRINE HYDROCHLORIDE, CYCLOBENZAPRINE HYDROCHLORIDE
 ERGOCALCIFEROL, ERGOCALCIFEROL
 FENOFIBRATE, FENOFIBRATE
 GLYCOPYRROLATE, GLYCOPYRROLATE
 LEVETIRACETAM, LEVETIRACETAM
 METRONIDAZOLE, METRONIDAZOLE

ORPHAN EUROPE

* ORPHAN EUROPE SARL

CYSTADANE, BETAINE

OSI PHARMS

* OSI PHARMACEUTICALS LLC

TARCEVA, ERLOTINIB HYDROCHLORIDE

OSMOTICA

* OSMOTICA KERESKEDELMI ES SZOLGALTATO KFT

HYDROMORPHONE HYDROCHLORIDE, HYDROMORPHONE HYDROCHLORIDE
 METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE

OSMOTICA PHARM

* OSMOTICA PHARMACEUTICAL

OSMOLEX ER, AMANTADINE HYDROCHLORIDE

* OSMOTICA PHARMACEUTICAL CORP

VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE

OSMOTICA PHARM US

* OSMOTICA PHARMACEUTICAL US LLC

NIFEDIPINE, NIFEDIPINE
 OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE

OTONOMY INC

* OTONOMY INC

OTIPRIO, CIPROFLOXACIN

OTSUKA

* OTSUKA PHARMACEUTICAL CO LTD

ABILIFY, ARIPIPRAZOLE

OTSUKA AMERICA PHARM

* OTSUKA AMERICA PHARMACEUTICAL INC

SAMSCA, TOLVAPTAN

OTSUKA PHARM

* OTSUKA PHARMACEUTICAL CO LTD

BUSULFEX, BUSULFAN

OTSUKA PHARM CO LTD

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** O ****

* OTSUKA PHARMACEUTICAL CO LTD
 ABILIFY MAINTENA KIT, ARIPIPIRAZOLE
 ABILIFY MYCITE KIT, ARIPIPIRAZOLE
 DACOGEN, DECITABINE
 JYNARQUE, TOLVAPTAN
 REXULTI, BREXPIPIRAZOLE

OUTLOOK PHARMS

* OUTLOOK PHARMACEUTICALS INC
 DEXTROAMPHETAMINE SULFATE, DEXTROAMPHETAMINE SULFATE

OXFORD PHARMS

* OXFORD PHARMACEUTICALS LLC
 ALPRAZOLAM, ALPRAZOLAM
 AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
 BACLOFEN, BACLOFEN
 BUSPIRONE HYDROCHLORIDE, BUSPIRONE HYDROCHLORIDE
 CYCLOBENZAPRINE HYDROCHLORIDE, CYCLOBENZAPRINE HYDROCHLORIDE
 GLYCOPYRROLATE, GLYCOPYRROLATE
 HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 LEVETIRACETAM, LEVETIRACETAM
 LORAZEPAM, LORAZEPAM
 METHOCARBAMOL, METHOCARBAMOL
 METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
 PAROXETINE HYDROCHLORIDE, PAROXETINE HYDROCHLORIDE
 PRIMIDONE, PRIMIDONE
 RIMACTANE, RIFAMPIN
 RISPERIDONE, RISPERIDONE
 SERTRALINE HYDROCHLORIDE, SERTRALINE HYDROCHLORIDE
 SIMVASTATIN, SIMVASTATIN
 SOTALOL HYDROCHLORIDE, SOTALOL HYDROCHLORIDE
 SPIRONOLACTONE, SPIRONOLACTONE
 TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE
 TRAZODONE HYDROCHLORIDE, TRAZODONE HYDROCHLORIDE

**** P ******P AND L**

* P AND L DEVELOPMENT LLC
 ADAPALENE, ADAPALENE
 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)
 M-ZOLE 3 COMBINATION PACK, MICONAZOLE NITRATE (OTC)
 MICONAZOLE 7, MICONAZOLE NITRATE (OTC)
 MICONAZOLE NITRATE, 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)

P AND L DEV LLC

* P AND L DEVELOPMENT LLC DBA PLD DEVELOPMENTS LLC
 IBUPROFEN, IBUPROFEN (OTC)

PACIFIC PHARMA

* PACIFIC PHARMA
 TIMOLOL MALEATE, TIMOLOL MALEATE
 * PACIFIC PHARMA INC
 TIMOLOL MALEATE, TIMOLOL MALEATE

PACIRA PHARMS INC

* PACIRA PHARMACEUTICALS INC
 EXPAREL, BUPIVACAINE

PADDOCK LLC

* PADDOCK LABORATORIES LLC
 ATOVAQUONE, ATOVAQUONE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** P ****

* PADDOCK LABORATORIES LLC
 BROMOCRIPTINE MESYLATE, BROMOCRIPTINE MESYLATE
 BROMPHENIRAMINE MALEATE, PSEUDOEPHEDRINE HYDROCHLORIDE AND DEXTROMETHORPHAN
 CALCIUM ACETATE, CALCIUM ACETATE
 CICLOPIROX, CICLOPIROX
 CLINDA-DERM, CLINDAMYCIN PHOSPHATE
 CLINDAMYCIN PALMITATE HYDROCHLORIDE, CLINDAMYCIN PALMITATE HYDROCHLORIDE
 CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
 CLOTRIMAZOLE, CLOTRIMAZOLE
 COLOCORT, HYDROCORTISONE
 COMPRO, PROCHLORPERAZINE
 DIHYDROERGOTAMINE MESYLATE, DIHYDROERGOTAMINE MESYLATE
 FLAVOXATE HYDROCHLORIDE, FLAVOXATE HYDROCHLORIDE
 HYDROCODONE BITARTRATE AND HOMATROPINE METHYLBROMIDE, HOMATROPINE METHYLBROMIDE
 HYDROCODONE BITARTRATE, CHLORPHENIRAMINE MALEATE AND PSEUDOEPHEDRINE HYDROCHLORIDE,
 HYDROMORPHONE HYDROCHLORIDE, HYDROMORPHONE HYDROCHLORIDE
 KIONEX, SODIUM POLYSTYRENE SULFONATE
 LAX-LYTE WITH FLAVOR PACKS, POLYETHYLENE GLYCOL 3350
 MIDAMOR, AMILORIDE HYDROCHLORIDE
 MIDAZOLAM HYDROCHLORIDE, MIDAZOLAM HYDROCHLORIDE
 MORPHINE SULFATE, MORPHINE SULFATE
 NARATRIPTAN, NARATRIPTAN HYDROCHLORIDE
 NYSTOP, NYSTATIN
 PODOFILOX, PODOFILOX
 POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
 REPAGLINIDE, REPAGLINIDE
 SODIUM POLYSTYRENE SULFONATE, SODIUM POLYSTYRENE SULFONATE
 TESTOSTERONE CYPIONATE, TESTOSTERONE CYPIONATE
 TROSPIUM CHLORIDE, TROSPIUM CHLORIDE

PANACEA BIOTEC LTD

* PANACEA BIOTEC LTD
 PRASUGREL, PRASUGREL HYDROCHLORIDE
 RIZATRIPTAN BENZOATE, RIZATRIPTAN BENZOATE
 TACROLIMUS, TACROLIMUS

PAR FORM

* PAR FORMULATIONS PRIVATE LTD
 DICLOFENAC POTASSIUM, DICLOFENAC POTASSIUM
 FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 LABETALOL HYDROCHLORIDE, LABETALOL HYDROCHLORIDE
 MAFENIDE ACETATE, MAFENIDE ACETATE
 ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE

PAR PHARM

* PAR PHARMACEUTICAL
 EVEROLIMUS, EVEROLIMUS
 PROPAFENONE HYDROCHLORIDE, PROPAFENONE HYDROCHLORIDE
 TESTOSTERONE, TESTOSTERONE

* PAR PHARMACEUTICAL INC
 ALPRAZOLAM, ALPRAZOLAM
 AMILORIDE HYDROCHLORIDE, AMILORIDE HYDROCHLORIDE
 AMLODIPINE BESYLATE AND BENAZEPRIL HYDROCHLORIDE, AMLODIPINE BESYLATE
 AMLODIPINE BESYLATE, VALSARTAN AND HYDROCHLOROTHIAZIDE, AMLODIPINE BESYLATE
 CABERGOLINE, CABERGOLINE
 CALCITONIN-SALMON, CALCITONIN SALMON
 CHOLESTYRAMINE LIGHT, CHOLESTYRAMINE
 CHOLESTYRAMINE, CHOLESTYRAMINE
 CLOMIPHENE CITRATE, CLOMIPHENE CITRATE
 CLONAZEPAM, CLONAZEPAM
 CYPROHEPTADINE HYDROCHLORIDE, CYPROHEPTADINE HYDROCHLORIDE
 DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
 DIPHENOXYLATE HYDROCHLORIDE AND ATROPINE SULFATE, ATROPINE SULFATE
 DOXEPIN HYDROCHLORIDE, DOXEPIN HYDROCHLORIDE
 DOXYCYCLINE, DOXYCYCLINE
 ESTAZOLAM, ESTAZOLAM
 FLUTAMIDE, FLUTAMIDE
 GLIPIZIDE, GLIPIZIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** P ******* PAR PHARMACEUTICAL INC**

GLYCOPYRROLATE, GLYCOPYRROLATE
 HYDRA-ZIDE, HYDRALAZINE HYDROCHLORIDE
 HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
 HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
 HYDROXYUREA, HYDROXYUREA
 IBUPROFEN, IBUPROFEN (OTC)
 IMIPRAMINE HYDROCHLORIDE, IMIPRAMINE HYDROCHLORIDE
 ISOSORBIDE DINITRATE, ISOSORBIDE DINITRATE
 LAMOTRIGINE, LAMOTRIGINE
 MEGESTROL ACETATE, MEGESTROL ACETATE
 METRONIDAZOLE, METRONIDAZOLE
 MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE
 MINOXIDIL, MINOXIDIL
 NATEGLINIDE, NATEGLINIDE
 NIFEDIPINE, NIFEDIPINE
 OLANZAPINE AND FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 OLANZAPINE, OLANZAPINE
 OMEPRAZOLE AND SODIUM BICARBONATE, OMEPRAZOLE
 OMEPRAZOLE AND SODIUM BICARBONATE, OMEPRAZOLE (OTC)
 OXANDROLONE, OXANDROLONE
 PIMOZIDE, PIMOZIDE
 POLYETHYLENE GLYCOL 3350, POLYETHYLENE GLYCOL 3350 (OTC)
 QUETIAPINE FUMARATE, QUETIAPINE FUMARATE
 RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
 RISPERIDONE, RISPERIDONE
 SODIUM PHENYLBUTYRATE, SODIUM PHENYLBUTYRATE
 SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
 TORSEMIDE, TORSEMIDE
 TRANLYCYPROMINE SULFATE, TRANLYCYPROMINE SULFATE
 TRAVOPROST, TRAVOPROST
 URSODIOL, URSODIOL
 VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE

PAR PHARM INC*** PAR PHARMACEUTICAL INC**

ACCOLATE, ZAFIRLUKAST
 ALOSETRON HYDROCHLORIDE, ALOSETRON HYDROCHLORIDE
 AMBRISENTAN, AMBRISENTAN
 AMLODIPINE BESYLATE AND VALSARTAN, AMLODIPINE BESYLATE
 ASPIRIN AND DIPYRIDAMOLE, ASPIRIN
 BOSENTAN, BOSENTAN
 DEXLANSOPRAZOLE, DEXLANSOPRAZOLE
 DEXMETHYLPHENIDATE HYDROCHLORIDE, DEXMETHYLPHENIDATE HYDROCHLORIDE
 DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE
 DOFETILIDE, DOFETILIDE
 DOXYLAMINE SUCCINATE AND PYRIDOXINE HYDROCHLORIDE, DOXYLAMINE SUCCINATE
 ERYTHROMYCIN ETHYLSUCCINATE, ERYTHROMYCIN ETHYLSUCCINATE
 ETHACRYNIC ACID, ETHACRYNIC ACID
 FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 ITRACONAZOLE, ITRACONAZOLE
 METOCLOPRAMIDE HYDROCHLORIDE, METOCLOPRAMIDE HYDROCHLORIDE
 MIDODRINE HYDROCHLORIDE, MIDODRINE HYDROCHLORIDE
 MORPHINE SULFATE, MORPHINE SULFATE
 OLMESARTAN MEDOXOMIL, AMLODIPINE AND HYDROCHLOROTHIAZIDE, AMLODIPINE BESYLATE
 PENICILLAMINE, PENICILLAMINE
 PHENOXYBENZAMINE HYDROCHLORIDE, PHENOXYBENZAMINE HYDROCHLORIDE
 PRAZIQUANTEL, PRAZIQUANTEL
 SAPROPTERIN DIHYDROCHLORIDE, SAPROPTERIN DIHYDROCHLORIDE
 TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE
 TOLCAPONE, TOLCAPONE
 TRIENTINE HYDROCHLORIDE, TRIENTINE HYDROCHLORIDE
 VIGABATRIN, VIGABATRIN
 ZILEUTON, ZILEUTON
 ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE

PAR STERILE PRODUCTS

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** P ****

* PAR STERILE PRODUCTS LLC
 ADRENALIN, EPINEPHRINE
 ARGATROBAN, ARGATROBAN
 BREVITAL SODIUM, METHOHEXITAL SODIUM
 BUPRENORPHINE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE
 COLY-MYCIN M, COLISTIMETHATE SODIUM
 CORPHEDRA, EPHEDRINE SULFATE
 DANTRIUM, DANTROLENE SODIUM
 DELESTROGEN, ESTRADIOL VALERATE
 DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE
 ETHACRYNATE SODIUM, ETHACRYNATE SODIUM
 FLUPHENAZINE DECANOATE, FLUPHENAZINE DECANOATE
 GANCICLOVIR SODIUM, GANCICLOVIR SODIUM
 KETALAR, KETAMINE HYDROCHLORIDE
 MYCOPHENOLATE MOFETIL HYDROCHLORIDE, MYCOPHENOLATE MOFETIL HYDROCHLORIDE
 NALOXONE HYDROCHLORIDE, NALOXONE HYDROCHLORIDE
 NEOSTIGMINE METHYLSULFATE, NEOSTIGMINE METHYLSULFATE
 PHENYLEPHRINE HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE
 PITOCIN, OXYTOCIN
 TIGAN, TRIMETHOBENZAMIDE HYDROCHLORIDE
 TREPROSTINIL, TREPROSTINIL
 TRIOSTAT, LIOTHYRONINE SODIUM
 VASOSTRICT, VASOPRESSIN

PARAGON BIOTECK

* PARAGON BIOTECK INC
 PHENYLEPHRINE HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE

PARAPRO LLC

* PARAPRO LLC
 NATROBA, SPINOSAD

PARATEK PHARMS INC

* PARATEK PHARMACEUTICALS INC
 NUZYRA, OMADACYCLINE TOSYLATE

PARKE DAVIS

* PARKE DAVIS DIV WARNER LAMBERT CO
 CELONTIN, METHSUXIMIDE
 CEREBYX, FOSPHENYTOIN SODIUM
 DILANTIN-125, PHENYTOIN
 NARDIL, PHENELZINE SULFATE
 NEURONTIN, GABAPENTIN
 ZARONTIN, ETHOSUXIMIDE

PARKE-DAVIS

* PARKE-DAVIS DIVISION OF PFIZER INC
 DILANTIN, PHENYTOIN SODIUM
 ZARONTIN, ETHOSUXIMIDE

PERRIGO

* PERRIGO CO
 MICONAZOLE NITRATE COMBINATION PACK, MICONAZOLE NITRATE (OTC)
 * PERRIGO LLC
 DESLORATADINE, DESLORATADINE

PERRIGO CO

* PERRIGO CO OF TENNESSEE INC
 CICLOPIROX, CICLOPIROX
 CLINDETS, CLINDAMYCIN PHOSPHATE
 CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
 ERYTHROMYCIN, ERYTHROMYCIN
 STIE-CORT, HYDROCORTISONE

PERRIGO CO TENNESSEE

* PERRIGO CO TENNESSEE INC
 BACITRACIN ZINC AND POLYMYXIN B SULFATE, BACITRACIN ZINC
 BACITRACIN, BACITRACIN
 BACITRACIN-NEOMYCIN-POLYMYXIN W/ HYDROCORTISONE ACETATE, BACITRACIN ZINC
 ERYTHROMYCIN, ERYTHROMYCIN
 GENTAMICIN SULFATE, GENTAMICIN SULFATE
 NEOMYCIN AND POLYMYXIN B SULFATES AND BACITRACIN ZINC, BACITRACIN ZINC
 NEOMYCIN AND POLYMYXIN B SULFATES AND DEXAMETHASONE, DEXAMETHASONE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** P ****

* PERRIGO CO TENNESSEE INC
SULFACETAMIDE SODIUM, SULFACETAMIDE SODIUM

PERRIGO ISRAEL

* PERRIGO ISRAEL PHARMACEUTICALS LTD
AMMONIUM LACTATE, AMMONIUM LACTATE
AZELASTINE HYDROCHLORIDE, AZELASTINE HYDROCHLORIDE
BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
CICLOPIROX, CICLOPIROX
CLINDAMYCIN PHOSPHATE AND BENZOYL PEROXIDE, BENZOYL PEROXIDE
CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
DESOXIMETASONE, DESOXIMETASONE
ECONAZOLE NITRATE, ECONAZOLE NITRATE
FLUOCINOLONE ACETONIDE, FLUOCINOLONE ACETONIDE
FLUOCINONIDE, FLUOCINONIDE
FLUTICASONE PROPIONATE, FLUTICASONE PROPIONATE
GYNAZOLE-1, BUTOCONAZOLE NITRATE
HALOBETASOL PROPIONATE, HALOBETASOL PROPIONATE
HYDROCORTISONE VALERATE, HYDROCORTISONE VALERATE
IMIQUIMOD, IMIQUIMOD
KETOCONAZOLE, KETOCONAZOLE
MESALAMINE, MESALAMINE
MINOXIDIL, MINOXIDIL (OTC)
MOMETASONE FUROATE, MOMETASONE FUROATE
MUPIROCIN, MUPIROCIN
NITROGLYCERIN, NITROGLYCERIN
OLOPATADINE HYDROCHLORIDE, OLOPATADINE HYDROCHLORIDE
PERMETHRIN, PERMETHRIN
PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
TERCONAZOLE, TERCONAZOLE
TESTOSTERONE, TESTOSTERONE
TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE (OTC)

PERRIGO NEW YORK

* PERRIGO NEW YORK INC
ACETAMINOPHEN, ACETAMINOPHEN (OTC)
BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
CENTANY, MUPIROCIN
CICLOPIROX, CICLOPIROX
CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
DESONIDE, DESONIDE
DESOXIMETASONE, DESOXIMETASONE
ERYTHROMYCIN, ERYTHROMYCIN
FLUTICASONE PROPIONATE, FLUTICASONE PROPIONATE
GENTAMICIN SULFATE, GENTAMICIN SULFATE
HYDROCORTISONE, HYDROCORTISONE
MINOXIDIL EXTRA STRENGTH (FOR MEN), MINOXIDIL (OTC)
MOMETASONE FUROATE, MOMETASONE FUROATE
NYSTATIN, NYSTATIN
PERMETHRIN, PERMETHRIN (OTC)
SELENIUM SULFIDE, SELENIUM SULFIDE
TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE

PERRIGO PHARMA INTL

* PERRIGO PHARMA INTERNATIONAL DAC
CLINDESSE, CLINDAMYCIN PHOSPHATE
ENTOCORT EC, BUDESONIDE
EVAMIST, ESTRADIOL
LORATADINE AND PSEUDOEPHEDRINE SULFATE, LORATADINE (OTC)
LORATADINE, LORATADINE (OTC)
PILOCARPINE HYDROCHLORIDE, PILOCARPINE HYDROCHLORIDE
TRETINOIN, TRETINOIN
* PERRIGO PHARMA INTERNATIONAL DESIGNATED ACTIVITY CO
LORATADINE, LORATADINE (OTC)
PREVACID 24 HR, LANSOPRAZOLE (OTC)

PERRIGO PHARMS CO

* PERRIGO PHARMACEUTICALS CO
ALBUTEROL SULFATE, ALBUTEROL SULFATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** P ****

* PERRIGO PHARMACEUTICALS CO
 CETIRIZINE HYDROCHLORIDE, CETIRIZINE HYDROCHLORIDE
 IBUPROFEN, IBUPROFEN
 NAPROXEN, NAPROXEN
 SCOPOLAMINE, SCOPOLAMINE

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
 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 SODIUM, IBUPROFEN SODIUM (OTC)
 IBUPROFEN, IBUPROFEN (OTC)
 LANSOPRAZOLE, LANSOPRAZOLE (OTC)
 LEVOCETIRIZINE DIHYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE (OTC)
 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)
 OMEPRAZOLE MAGNESIUM, OMEPRAZOLE MAGNESIUM (OTC)
 POLYETHYLENE GLYCOL 3350, POLYETHYLENE GLYCOL 3350 (OTC)
 RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE (OTC)

PERRIGO UK FINCO

* PERRIGO UK FINCO LTD PARTNERSHIP
 ACYCLOVIR, ACYCLOVIR
 ADAPALENE AND BENZOYL PEROXIDE, ADAPALENE
 BETAMETHASONE VALERATE, BETAMETHASONE VALERATE
 CALCIPOTRIENE AND BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
 CLINDAMYCIN PHOSPHATE AND BENZOYL PEROXIDE, BENZOYL PEROXIDE
 CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
 CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
 DICLOFENAC SODIUM, DICLOFENAC SODIUM
 ESTRADIOL, ESTRADIOL
 FLURANDRENOLIDE, FLURANDRENOLIDE
 INGENOL MEBUTATE, INGENOL MEBUTATE
 METRONIDAZOLE, METRONIDAZOLE
 NYSTATIN AND TRIAMCINOLONE ACETONIDE, NYSTATIN
 TESTOSTERONE, TESTOSTERONE
 TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE

PERSION

* PERSION PHARMACEUTICALS LLC
 ZOHYDRO ER, HYDROCODONE BITARTRATE

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
 CHANTIX, VARENICLINE TARTRATE
 INLYTA, AXITINIB
 LYRICA CR, PREGABALIN
 LYRICA, PREGABALIN

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** P ****

* PF PRISM CV

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

CADUET, AMLODIPINE BESYLATE
 CALAN SR, VERAPAMIL HYDROCHLORIDE
 CARDURA XL, DOXAZOSIN MESYLATE
 ELELYSO, TALIGLUCERASE ALFA
 GEODON, ZIPRASIDONE HYDROCHLORIDE
 GEODON, ZIPRASIDONE MESYLATE
 GLUCOTROL XL, GLIPIZIDE
 GLUCOTROL, GLIPIZIDE
 HEPARIN SODIUM PRESERVATIVE FREE, HEPARIN SODIUM
 HEPARIN SODIUM, HEPARIN SODIUM
 LIPITOR, ATORVASTATIN CALCIUM
 MERREM, MEROPENEM
 NORVASC, AMLODIPINE BESYLATE
 PROCARDIA, NIFEDIPINE
 REVATIO, SILDENAFIL CITRATE
 SONATA, ZALEPLON
 TESSALON, BENZONATATE
 TOVIAZ, FESOTERODINE FUMARATE
 UNASYN, AMPICILLIN SODIUM
 ZITHROMAX, AZITHROMYCIN

* PFIZER LABORATORIES DIV PFIZER INC

CARDURA, DOXAZOSIN MESYLATE
 FELDENE, PIROXICAM
 MINIPRESS, PRAZOSIN HYDROCHLORIDE
 PFIZERPEN, PENICILLIN G POTASSIUM
 PROCARDIA XL, NIFEDIPINE
 UNASYN, AMPICILLIN SODIUM
 VIBRAMYCIN, DOXYCYCLINE
 VIBRAMYCIN, DOXYCYCLINE CALCIUM
 VIBRAMYCIN, DOXYCYCLINE HYCLATE

* PFIZER PHARMACEUTICALS INC

DILANTIN, PHENYTOIN
 ZOLOFT, SERTRALINE HYDROCHLORIDE

* PFIZER PHARMACEUTICALS PRODUCTION CORP LTD

TIKOSYN, DOFETILIDE

PFIZER INC

* PFIZER INC

ADVIL DUAL ACTION WITH ACETAMINOPHEN, ACETAMINOPHEN (OTC)
 CAMPTOSAR, IRINOTECAN HYDROCHLORIDE
 DAURISMO, GLASDEGIB MALEATE
 ELLENCE, EPIRUBICIN HYDROCHLORIDE
 FRAGMIN, DALTEPARIN SODIUM
 IBRANCE, PALBOCICLIB
 LORBRENA, LORLATINIB
 NICOTROL, NICOTINE
 TALZENNA, TALAZOPARIB TOSYLATE
 VIAGRA, SILDENAFIL CITRATE
 VIZIMPRO, DACOMITINIB
 XELJANZ XR, TOFACITINIB CITRATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** P ******PFIZER IRELAND**

* PFIZER IRELAND PHARMACEUTICALS
RELPAF, ELETRIPTAN HYDROBROMIDE

PFIZER PHARMS

* PFIZER PHARMACEUTICALS LTD
ACCUPRIL, QUINAPRIL HYDROCHLORIDE
ACCURETIC, HYDROCHLOROTHIAZIDE
LOPID, GEMFIBROZIL
NEURONTIN, GABAPENTIN
NITROSTAT, NITROGLYCERIN

PHARM ASSOC

* PHARMACEUTICAL ASSOC INC
CYPROHEPTADINE HYDROCHLORIDE, CYPROHEPTADINE HYDROCHLORIDE
HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
LORAZEPAM, LORAZEPAM
PREDNISOLONE SODIUM PHOSPHATE, PREDNISOLONE SODIUM PHOSPHATE

* PHARMACEUTICAL ASSOCIATES INC
ACETAMINOPHEN AND CODEINE PHOSPHATE, ACETAMINOPHEN
AMANTADINE HYDROCHLORIDE, AMANTADINE HYDROCHLORIDE
CHLORHEXIDINE GLUCONATE, CHLORHEXIDINE GLUCONATE
CIMETIDINE HYDROCHLORIDE, CIMETIDINE HYDROCHLORIDE
DIPHENHYDRAMINE HYDROCHLORIDE, DIPHENHYDRAMINE HYDROCHLORIDE
ESCITALOPRAM OXALATE, ESCITALOPRAM OXALATE
ETHOSUXIMIDE, ETHOSUXIMIDE
FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
FLUPHENAZINE HYDROCHLORIDE, FLUPHENAZINE HYDROCHLORIDE
HALOPERIDOL, HALOPERIDOL LACTATE
HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
LACTULOSE, LACTULOSE
LEVETIRACETAM, LEVETIRACETAM
METHADONE HYDROCHLORIDE, METHADONE HYDROCHLORIDE
METOCLOPRAMIDE HYDROCHLORIDE, METOCLOPRAMIDE HYDROCHLORIDE
MORPHINE SULFATE, MORPHINE SULFATE
NORTRIPTYLINE HYDROCHLORIDE, NORTRIPTYLINE HYDROCHLORIDE
NYSTATIN, NYSTATIN
ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE
OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
PREDNISOLONE SODIUM PHOSPHATE, PREDNISOLONE SODIUM PHOSPHATE
PREDNISOLONE, PREDNISOLONE
RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
RISPERIDONE, RISPERIDONE
SULFATRIM PEDIATRIC, SULFAMETHOXAZOLE
THEOPHYLLINE, THEOPHYLLINE
TRIHEXYPHENIDYL HYDROCHLORIDE, TRIHEXYPHENIDYL HYDROCHLORIDE
VALPROIC ACID, VALPROIC ACID
VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE

PHARM SOURCING

* PHARMACEUTICAL SOURCING PARTNERS INC
MESALAMINE, MESALAMINE

PHARMA RES SOFTWARE

* PHARMA RESEARCH SOFTWARE SOLUTION LLC
POTASSIUM CHLORIDE, POTASSIUM CHLORIDE

PHARMACHEMIE BV

* PHARMACHEMIE BV
CARBOPLATIN, CARBOPLATIN
CISPLATIN, CISPLATIN
DOXORUBICIN HYDROCHLORIDE, DOXORUBICIN HYDROCHLORIDE
METHOTREXATE SODIUM PRESERVATIVE FREE, METHOTREXATE SODIUM

PHARMACIA

* PHARMACIA AND UPJOHN CO LLC
GENOTROPIN PRESERVATIVE FREE, SOMATROPIN
GENOTROPIN, SOMATROPIN
SOMAVERT, PEGVISOMANT

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** P ******PHARMACIA AND UPJOHN**

- * PHARMACIA AND UPJOHN
 - XANAX XR, ALPRAZOLAM
- * PHARMACIA AND UPJOHN CO
 - AROMASIN, EXEMESTANE
 - AZULFIDINE EN-TABS, SULFASALAZINE
 - AZULFIDINE, SULFASALAZINE
 - BACITRACIN, BACITRACIN
 - CAVERJECT IMPULSE, ALPROSTADIL
 - CAVERJECT, ALPROSTADIL
 - CLEOCIN HYDROCHLORIDE, CLINDAMYCIN HYDROCHLORIDE
 - CLEOCIN PHOSPHATE IN DEXTROSE 5% IN PLASTIC CONTAINER, CLINDAMYCIN PHOSPHATE
 - CLEOCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
 - CLEOCIN T, CLINDAMYCIN PHOSPHATE
 - CLEOCIN, CLINDAMYCIN PALMITATE HYDROCHLORIDE
 - CLEOCIN, CLINDAMYCIN PHOSPHATE
 - CORTEF, HYDROCORTISONE
 - CORVERT, IBUTILIDE FUMARATE
 - CYKLOKAPRON, TRANEXAMIC ACID
 - DEPO-ESTRADIOL, ESTRADIOL CYPIONATE
 - DEPO-MEDROL, METHYLPREDNISOLONE ACETATE
 - DEPO-PROVERA, MEDROXYPROGESTERONE ACETATE
 - DEPO-TESTOSTERONE, TESTOSTERONE CYPIONATE
 - DETROL LA, TOLTERODINE TARTRATE
 - DETROL, TOLTERODINE TARTRATE
 - DOXORUBICIN HYDROCHLORIDE, DOXORUBICIN HYDROCHLORIDE
 - EMCYT, ESTRAMUSTINE PHOSPHATE SODIUM
 - ESTRING, ESTRADIOL
 - GLYNASE, GLYBURIDE
 - GLYSET, MIGLITOL
 - HALCION, TRIAZOLAM
 - HEMABATE, CARBOPROST TROMETHAMINE
 - IDAMYCIN PFS, IDARUBICIN HYDROCHLORIDE
 - LINCOCIN, LINCOMYCIN HYDROCHLORIDE
 - MEDROL, METHYLPREDNISOLONE
 - MYCOBUTIN, RIFABUTIN
 - NICOTROL, NICOTINE
 - OGEN 5, ESTROPIPATE
 - PREPIDIL, DINOPROSTONE
 - PROSTIN E2, DINOPROSTONE
 - PROSTIN VR PEDIATRIC, ALPROSTADIL
 - PROVERA, MEDROXYPROGESTERONE ACETATE
 - R-GENE 10, ARGININE HYDROCHLORIDE
 - SOLU-CORTEF, HYDROCORTISONE SODIUM SUCCINATE
 - SOLU-MEDROL, METHYLPREDNISOLONE SODIUM SUCCINATE
 - XALATAN, LATANOPROST
 - XANAX, ALPRAZOLAM
 - ZINECARD, DEXRAZOXANE HYDROCHLORIDE
 - ZYVOX, LINEZOLID
- * PHARMACIA AND UPJOHN SUB PFIZER INC
 - DEPO-SUBQ PROVERA 104, MEDROXYPROGESTERONE ACETATE

PHARMACIA UPJOHN

- * PHARMACIA AND UPJOHN CO A SUB OF PFIZER INC
 - COLESTID, COLESTIPOL HYDROCHLORIDE
 - FLAVORED COLESTID, COLESTIPOL HYDROCHLORIDE

PHARMACOSMOS AS

- * PHARMACOSMOS AS
 - MONOFERRIC, FERRIC DERISOMALTOSE

PHARMACYCLICS INC

- * PHARMACYCLICS INC
 - IMBRUVICA, IBRUTINIB

PHARMADAX INC

- * PHARMADAX INC
 - GLYBURIDE, GLYBURIDE
 - LEVETIRACETAM, LEVETIRACETAM

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** P ****

* PHARMADAX INC

QUETIAPINE FUMARATE, QUETIAPINE FUMARATE

PHARMALUCENCE

* PHARMALUCENCE INC

AN-SULFUR COLLOID, TECHNETIUM TC-99M SULFUR COLLOID KIT

CIS-MDP, TECHNETIUM TC-99M MEDRONATE KIT

CIS-PYRO, TECHNETIUM TC-99M PYROPHOSPHATE KIT

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

PHARMASCIENCE INC

* PHARMASCIENCE INC

BUSULFAN, BUSULFAN

DECITABINE, DECITABINE

GANCICLOVIR SODIUM, GANCICLOVIR SODIUM

PHARMAXIS LTD

* PHARMAXIS LTD

ARIDOL KIT, MANNITOL

PHOTOCURE ASA

* PHOTOCURE ASA

CYSVIEW KIT, HEXAMINOLEVULINATE HYDROCHLORIDE

PIERRE FABRE DERMA

* PIERRE FABRE DERMATOLOGIE

HEMANGEOL, PROPRANOLOL HYDROCHLORIDE

PIERREL

* PIERREL S.P.A.

ORABLOC, ARTICAINA HYDROCHLORIDE

PII

* PHARMACEUTICS INTERNATIONAL INC

BUTALBITAL, ASPIRIN AND CAFFEINE, ASPIRIN

ITRACONAZOLE, ITRACONAZOLE

PIROXICAM, PIROXICAM

POTASSIUM CHLORIDE, POTASSIUM CHLORIDE

PINNACLE BIOLGS

* PINNACLE BIOLOGICS INC

PHOTOFRIN, PORFIMER SODIUM

PIRAMAL CRITICAL

* PIRAMAL CRITICAL CARE INC

GABLOFEN, BACLOFEN

GLYCOPYRROLATE, GLYCOPYRROLATE

ISOFLURANE, ISOFLURANE

MITIGO, MORPHINE SULFATE

OXACILLIN SODIUM, OXACILLIN SODIUM

SOJOURN, SEVOFLURANE

* PIRAMAL CRITICAL CARE LTD

LEVOTHYROXINE SODIUM, LEVOTHYROXINE SODIUM

PIRAMAL ENT

* PIRAMAL ENTERPRISES LTD

ISOFLURANE, ISOFLURANE

PIRAMAL HLTHCARE UK

* PIRAMAL HEALTHCARE UK LTD

CINACALCET HYDROCHLORIDE, CINACALCET HYDROCHLORIDE

CLOBAZAM, CLOBAZAM

DEFERASIROX, DEFERASIROX

TETRABENAZINE, TETRABENAZINE

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

LORATADINE, LORATADINE (OTC)

ZOLMITRIPTAN, ZOLMITRIPTAN

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** P ******PLIVA**

* PLIVA INC
 AZITHROMYCIN, AZITHROMYCIN
 BENZTROPINE MESYLATE, BENZTROPINE MESYLATE
 CIMETIDINE, CIMETIDINE
 CYCLOBENZAPRINE HYDROCHLORIDE, CYCLOBENZAPRINE HYDROCHLORIDE
 DOXAZOSIN MESYLATE, DOXAZOSIN MESYLATE
 HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
 HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE
 KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
 METRONIDAZOLE, METRONIDAZOLE
 NAPROXEN, NAPROXEN
 TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE
 TRAZODONE HYDROCHLORIDE, TRAZODONE HYDROCHLORIDE
 TRIAMTERENE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 WARFARIN SODIUM, WARFARIN SODIUM

PLIVA HRVATSKA DOO

* PLIVA HRVATSKA DOO
 DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
 ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE

PLIVA LACHEMA

* PLIVA LACHEMA AS
 CARBOPLATIN, CARBOPLATIN
 IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE

PLIVA PHARM IND

* PLIVA PHARMACEUTICAL INDUSTRY INC
 TORSEMIDE, TORSEMIDE

PLX PHARMA

* PLX PHARMA INC
 VAZALORE, ASPIRIN (OTC)

POHL BOSKAMP

* POHL BOSKAMP
 NITROLINGUAL PUMPSPRAY, NITROGLYCERIN

POLYGEN PHARMS

* POLYGEN PHARMACEUTICALS INC
 AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
 DARIFENACIN HYDROBROMIDE, DARIFENACIN HYDROBROMIDE
 PHENTERMINE HYDROCHLORIDE, PHENTERMINE HYDROCHLORIDE

POLYMEDICA

* POLYMEDICA INDUSTRIES INC
 NEOPAP, ACETAMINOPHEN (OTC)

PORTOLA PHARMS INC

* PORTOLA PHARMACEUTICALS INC
 BEVYXXA, BETRIXABAN

POWDER PHARMS

* POWDER PHARMACEUTICALS INC
 ZINGO, LIDOCAINE HYDROCHLORIDE

PRAGMA

* PRAGMA PHARMACEUTICALS LLC
 KEFLEX, CEPHALEXIN

PRAXAIR DISTRIBUTION

* PRAXAIR DISTRIBUTION INC
 NOXIVENT, NITRIC OXIDE

PRECISION DERMAT

* PRECISION DERMATOLOGY INC
 LOCID LIPOCREAM, HYDROCORTISONE BUTYRATE
 LOCID, 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

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** P ****

* PRECISION NUCLEAR LLC
SODIUM FLUORIDE F-18, SODIUM FLUORIDE F-18

PRINSTON INC

* PRINSTON PHARMACEUTICAL INC
AMPHETAMINE SULFATE, AMPHETAMINE SULFATE
ARIPIPIRAZOLE, ARIPIPIRAZOLE
BENZAEPRIIL HYDROCHLORIDE, BENAZEPRIL HYDROCHLORIDE
BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
CANDESARTAN CILEXETIL AND HYDROCHLOROTHIAZIDE, CANDESARTAN CILEXETIL
CAPTOPRIL, CAPTOPRIL
CLONAZEPAM, CLONAZEPAM
CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
CLOPIDOGREL BISULFATE, CLOPIDOGREL BISULFATE
CYCLOBENZAPRINE HYDROCHLORIDE, CYCLOBENZAPRINE HYDROCHLORIDE
DIVALPROEX SODIUM, DIVALPROEX SODIUM
DOFETILIDE, DOFETILIDE
DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE
ENTECAVIR, ENTECAVIR
ESCITALOPRAM OXALATE, ESCITALOPRAM OXALATE
FENOFIBRATE, FENOFIBRATE
FOSINOPRIL SODIUM, FOSINOPRIL SODIUM
FUROSEMIDE, FUROSEMIDE
GLIMEPIRIDE, GLIMEPIRIDE
GLYCOPYRROLATE, GLYCOPYRROLATE
HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE
IRBESARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
IRBESARTAN, IRBESARTAN
LEVETIRACETAM, LEVETIRACETAM
LISINOPRIL AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
LISINOPRIL, LISINOPRIL
LOSARTAN POTASSIUM AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
LOSARTAN POTASSIUM, LOSARTAN POTASSIUM
METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
METHOCARBAMOL, METHOCARBAMOL
NEVIRAPINE, NEVIRAPINE
OLMESARTAN MEDOXOMIL AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
PAROXETINE MESYLATE, PAROXETINE MESYLATE
PAROXETINE, PAROXETINE HYDROCHLORIDE
PHENTERMINE HYDROCHLORIDE, PHENTERMINE HYDROCHLORIDE
PIOGLITAZONE HYDROCHLORIDE, PIOGLITAZONE HYDROCHLORIDE
POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
QUINAPRIL HYDROCHLORIDE, QUINAPRIL HYDROCHLORIDE
RISPERIDONE, RISPERIDONE
ROPINIROLE HYDROCHLORIDE, ROPINIROLE HYDROCHLORIDE
TELMISARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
TELMISARTAN, TELMISARTAN
TEMAZEPAM, TEMAZEPAM
VALSARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
VALSARTAN, VALSARTAN
VORICONAZOLE, VORICONAZOLE

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
AZEDRA, IOBENGUANE I-131

PROVELL

* PROVELL PHARMACEUTICALS LLC
EUTHYROX, LEVOTHYROXINE SODIUM **

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** P ******PROVENSIS**

* PROVENSIS LTD
VARITHENA, POLIDOCANOL

PROVEPHARM SAS

* PROVEPHARM SAS
PROVAYBLUE, METHYLENE BLUE

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)
PIOGLITAZONE HYDROCHLORIDE, PIOGLITAZONE HYDROCHLORIDE

PURDUE

* PURDUE GMP CENTER LLC
SEROMYCIN, CYCLOSERINE

PURDUE PHARMA LP

* PURDUE PHARMA LP
ADHANSIA XR, METHYLPHENIDATE HYDROCHLORIDE
BUTRANS, BUPRENORPHINE
HYSINGLA ER, HYDROCODONE BITARTRATE
MS CONTIN, MORPHINE SULFATE
OXYCONTIN, OXYCODONE HYDROCHLORIDE

**** 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
CEFAZOLIN SODIUM, CEFAZOLIN SODIUM
CEFEPIME HYDROCHLORIDE, CEFEPIME HYDROCHLORIDE
CEFTRIAZONE, CEFTRIAZONE SODIUM
IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE
OLANZAPINE, OLANZAPINE
OLMESARTAN MEDOXOMIL, OLMESARTAN MEDOXOMIL
ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
OXALIPLATIN, OXALIPLATIN
PALONOSETRON HYDROCHLORIDE, PALONOSETRON HYDROCHLORIDE
PIPERACILLIN AND TAZOBACTAM, PIPERACILLIN SODIUM
SOLIFENACIN SUCCINATE, SOLIFENACIN SUCCINATE
TADALAFIL, TADALAFIL
TENOFIVIR DISOPROXIL FUMARATE, TENOFIVIR DISOPROXIL FUMARATE

QINGDAO BAHEAL PHARM

* QINGDAO BAHEAL PHARMACEUTICAL CO LTD
BENZONATATE, BENZONATATE
CELECOXIB, CELECOXIB
DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE
FOLIC ACID, FOLIC ACID

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** Q ****

* QINGDAO BAHEAL PHARMACEUTICAL CO LTD
METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE

QOL MEDCL

* QOL MEDICAL LLC
ETHAMOLIN, ETHANOLAMINE OLEATE
SUCRAID, SACROSIDASE

QUAGEN

* QUAGEN PHARMACEUTICALS LLC
ALBUTEROL SULFATE, ALBUTEROL SULFATE
CHILDREN'S CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
CYPROHEPTADINE HYDROCHLORIDE, CYPROHEPTADINE HYDROCHLORIDE
PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE

QUEEN HAMAMATSU PET

* QUEEN HAMAMATSU PET IMAGING CENTER
FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18

**** R ******R-PHARM US LLC**

* R-PHARM US LLC
IXEMPRA KIT, IXABEPILONE

RADIUS HEALTH INC

* RADIUS HEALTH INC
TYMLOS, ABALOPARATIDE

RANBAXY

* RANBAXY SIGNATURE LLC
RIOMET, METFORMIN HYDROCHLORIDE

RB HLTH

* RB HEALTH US LLC
DELSYM, DEXTROMETHORPHAN POLISTIREX (OTC)
MUCINEX D, GUAIFENESIN (OTC)
MUCINEX DM, DEXTROMETHORPHAN HYDROBROMIDE (OTC)
MUCINEX, GUAIFENESIN (OTC)

RECIP

* RECIP AB
THYROSAFE, POTASSIUM IODIDE (OTC)

RECIPHARM

* RECIPHARM PHARMASERVICES PRIVATE LTD
FLUCYTOSINE, FLUCYTOSINE

RECORDATI RARE

* RECORDATI RARE DISEASES INC
CARBAGLU, CARGLUMIC ACID
CHEMET, SUCCIMER
COSMEGEN, DACTINOMYCIN
DESOXYN, METHAMPHETAMINE HYDROCHLORIDE
INDOCIN, INDOMETHACIN SODIUM
NEOPROFEN, IBUPROFEN LYSINE
PEGANONE, ETHOTOIN
SIGNIFOR LAR KIT, PASIREOTIDE PAMOATE
SIGNIFOR, PASIREOTIDE DIASPARTATE
TRANXENE, CLORAZEPATE DIPOTASSIUM

RECRO GAINESVILLE

* RECRO GAINESVILLE LLC
VERELAN PM, VERAPAMIL HYDROCHLORIDE
VERELAN, VERAPAMIL HYDROCHLORIDE

REDHILL

* REDHILL BIOPHARMA INC
AEMCOLO, RIFAMYCIN SODIUM
* REDHILL BIOPHARMA LTD
TALICIA, AMOXICILLIN

RELYPSA INC

* RELYPSA INC
VELTASSA, PATIROMER SORBITEX CALCIUM

REMPEX PHARMS

* REMPEX PHARMACEUTICALS

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** R ****

- * REMPEX PHARMACEUTICALS
MINOCIN, MINOCYCLINE HYDROCHLORIDE
- * REMPEX PHARMACEUTICALS A WHOLLY OWNED SUB OF MELINTA THERAPEUTICS INC
VABOMERE, MEROPENEM

RENATA

- * RENATA LTD
RISPERIDONE, RISPERIDONE

RHODES PHARMS

- * RHODES PHARMACEUTICALS LP
APTENSIO XR, METHYLPHENIDATE 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
MORPHINE SULFATE, MORPHINE SULFATE
OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN
OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
THEOPHYLLINE, THEOPHYLLINE

RICONPHARMA LLC

- * RICONPHARMA LLC
BETAMETHASONE VALERATE, BETAMETHASONE VALERATE
CHLORTHALIDONE, CHLORTHALIDONE
DOFETILIDE, DOFETILIDE
GLYCOPYRROLATE, GLYCOPYRROLATE
ISOSORBIDE MONONITRATE, ISOSORBIDE MONONITRATE
PHENYLEPHRINE HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE

RIGEL PHARMS INC

- * RIGEL PHARMACEUTICALS INC
TAVALISSE, FOSTAMATINIB DISODIUM

RISING

- * RISING PHARMA HOLDINGS INC
ABIRATERONE ACETATE, ABIRATERONE ACETATE
ACETIC ACID, ACETIC ACID, GLACIAL
ALOSETRON HYDROCHLORIDE, ALOSETRON HYDROCHLORIDE
BUMETANIDE, BUMETANIDE
CEVIMELINE HYDROCHLORIDE, CEVIMELINE HYDROCHLORIDE
CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE
CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
CROMOLYN SODIUM, CROMOLYN SODIUM
DESOXIMETASONE, DESOXIMETASONE
DICLOFENAC SODIUM, DICLOFENAC SODIUM
DIFLORASONE DIACETATE, DIFLORASONE DIACETATE
DOXERCALCIFEROL, DOXERCALCIFEROL
GLYCOPYRROLATE, GLYCOPYRROLATE
HYDROCORTISONE, HYDROCORTISONE
LANSOPRAZOLE, AMOXICILLIN AND CLARITHROMYCIN, AMOXICILLIN
LEVOCARNITINE, LEVOCARNITINE
LEVOFLOXACIN, LEVOFLOXACIN
METHIMAZOLE, METHIMAZOLE
METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
NYSTATIN AND TRIAMCINOLONE ACETONIDE, NYSTATIN
PARICALCITOL, PARICALCITOL
PERPHENAZINE, PERPHENAZINE
PREGABALIN, PREGABALIN
PYRIDOSTIGMINE BROMIDE, PYRIDOSTIGMINE BROMIDE
RISPERIDONE, RISPERIDONE
SELEGILINE HYDROCHLORIDE, SELEGILINE HYDROCHLORIDE
TEMOZOLOMIDE, TEMOZOLOMIDE
TOREMIFENE CITRATE, TOREMIFENE CITRATE
TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE
TRIENTINE HYDROCHLORIDE, TRIENTINE HYDROCHLORIDE
ZILEUTON, ZILEUTON

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** R ****

* RISING PHARMACEUTICALS
URSODIOL, URSODIOL

ROCHE PALO

* ROCHE PALO ALTO LLC
CELLCEPT, MYCOPHENOLATE MOFETIL
CELLCEPT, MYCOPHENOLATE MOFETIL HYDROCHLORIDE

ROCKWELL MEDICAL INC

* ROCKWELL MEDICAL INC
TRIFERIC, FERRIC PYROPHOSPHATE CITRATE

ROMARK

* ROMARK LABORATORIES
ALINIA, NITAZOXANIDE

ROUSES POINT PHARMS

* ROUSES POINT PHARMACEUTICALS LLC
LEVETIRACETAM, LEVETIRACETAM

RTRX

* RETROPHIN INC
CHOLBAM, CHOLIC ACID

RUBICON

* RUBICON RESEARCH PRIVATE LTD
AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE
BACLOFEN, BACLOFEN
CARVEDILOL, CARVEDILOL
CYCLOBENZAPRINE HYDROCHLORIDE, CYCLOBENZAPRINE HYDROCHLORIDE
GABAPENTIN, GABAPENTIN
LAMOTRIGINE, LAMOTRIGINE
METOPROLOL TARTRATE, METOPROLOL TARTRATE
MIDODRINE HYDROCHLORIDE, MIDODRINE HYDROCHLORIDE
OXCARBAZEPINE, OXCARBAZEPINE
PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
RABEPRAZOLE SODIUM, RABEPRAZOLE SODIUM
SILDENAFIL CITRATE, SILDENAFIL CITRATE
TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE

RXMTM THERAPS LLC

* RXMTM THERAPEUTICS LLC A WHOLLY OWNED SUB OF CUTISPHARMA INC
FIRVANQ KIT, VANCOMYCIN HYDROCHLORIDE

**** S ******SAGE PRODS**

* SAGE PRODUCTS INC
CHLORHEXIDINE GLUCONATE, CHLORHEXIDINE GLUCONATE (OTC)

SAGE THERAP

* SAGE THERAPEUTICS INC
ZULRESSO, BREXANOLONE

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

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** S ******* SAGENT PHARMACEUTICALS INC**

CEFEPIME HYDROCHLORIDE, CEFEPIME HYDROCHLORIDE
 CHLOROTHIAZIDE SODIUM, CHLOROTHIAZIDE SODIUM
 CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
 COLISTIMETHATE SODIUM, COLISTIMETHATE SODIUM
 DAPTOMYCIN, DAPTOMYCIN
 DECITABINE, DECITABINE
 DESMOPRESSIN ACETATE, DESMOPRESSIN ACETATE
 DIHYDROERGOTAMINE MESYLATE, DIHYDROERGOTAMINE MESYLATE
 EPTIFIBATIDE, EPTIFIBATIDE
 ESMOLOL HYDROCHLORIDE, ESMOLOL HYDROCHLORIDE
 FLUDARABINE PHOSPHATE, FLUDARABINE PHOSPHATE
 FLUOROURACIL, FLUOROURACIL
 FULVESTRANT, FULVESTRANT
 GEMCITABINE HYDROCHLORIDE, GEMCITABINE HYDROCHLORIDE
 GLYDO, LIDOCAINE HYDROCHLORIDE
 IBANDRONATE SODIUM, IBANDRONATE SODIUM
 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
 OCTREOTIDE ACETATE (PRESERVATIVE FREE), OCTREOTIDE ACETATE
 OCTREOTIDE ACETATE, OCTREOTIDE ACETATE
 OXYTOCIN, OXYTOCIN
 PALONOSETRON HYDROCHLORIDE, PALONOSETRON HYDROCHLORIDE
 PAMIDRONATE DISODIUM, PAMIDRONATE DISODIUM
 PENTOBARBITAL SODIUM, PENTOBARBITAL SODIUM
 PROPOFOL, PROPOFOL
 ROCURONIUM BROMIDE, ROCURONIUM BROMIDE
 SODIUM NITROPRUSSIDE, SODIUM NITROPRUSSIDE
 THIAMINE HYDROCHLORIDE, THIAMINE HYDROCHLORIDE
 VECURONIUM BROMIDE, VECURONIUM BROMIDE
 ZOLEDRONIC ACID, ZOLEDRONIC ACID

SAI LIFE SCIENCES*** SAI LIFE SCIENCES LTD**

DAPSONE, DAPSONE

SALIX*** SALIX PHARMACEUTICALS INC**

FENOGLIDE, FENOFIBRATE
 PLENVU, ASCORBIC ACID
 RELISTOR, METHYLNALTREXONE BROMIDE
 TRULANCE, PLECANATIDE
 UCERIS, BUDESONIDE
 ZEGERID, OMEPRAZOLE

SALIX PHARMS*** SALIX PHARMACEUTICALS INC**

ANUSOL HC, HYDROCORTISONE
 DIURIL, CHLOROTHIAZIDE
 MOVIPREP, ASCORBIC ACID
 OSMOPREP, SODIUM PHOSPHATE, DIBASIC, ANHYDROUS
 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

SANDOZ

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** S **

* SANDOZ
 DOCETAXEL, DOCETAXEL

* SANDOZ INC
 ALPRAZOLAM, ALPRAZOLAM
 AMANTADINE HYDROCHLORIDE, AMANTADINE HYDROCHLORIDE
 AMIODARONE HYDROCHLORIDE, AMIODARONE 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
 APREPITANT, APREPITANT
 ARGATROBAN IN SODIUM CHLORIDE, ARGATROBAN
 AZITHROMYCIN, AZITHROMYCIN
 BENAZEPRIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, BENAZEPRIL HYDROCHLORIDE
 BICALUTAMIDE, BICALUTAMIDE
 BISOPROLOL FUMARATE AND HYDROCHLOROTHIAZIDE, BISOPROLOL FUMARATE
 BUMETANIDE, BUMETANIDE
 BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
 CAFERGOT, CAFFEINE
 CARISOPRODOL AND ASPIRIN, ASPIRIN
 CARISOPRODOL, ASPIRIN AND CODEINE PHOSPHATE, ASPIRIN
 CARVEDILOL, CARVEDILOL
 CEFAZOLIN SODIUM, CEFAZOLIN SODIUM
 CEFDINIR, CEFDINIR
 CEFPODOXIME PROXETIL, CEFPODOXIME PROXETIL
 CEFPROZIL, CEFPROZIL
 CEFTRIAZONE, CEFTRIAZONE SODIUM
 CHILDREN'S CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 CHOLESTYRAMINE LIGHT, CHOLESTYRAMINE
 CHOLESTYRAMINE, CHOLESTYRAMINE
 CLARITHROMYCIN, CLARITHROMYCIN
 CLOMIPRAMINE HYDROCHLORIDE, CLOMIPRAMINE HYDROCHLORIDE
 CLONAZEPAM, CLONAZEPAM
 COSYNTROPIN, COSYNTROPIN
 CYCLOSPORINE, CYCLOSPORINE
 DESIPRAMINE HYDROCHLORIDE, DESIPRAMINE HYDROCHLORIDE
 DESLORATADINE, DESLORATADINE
 DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE
 DICLOFENAC SODIUM AND MISOPROSTOL, DICLOFENAC SODIUM
 DICLOXACILLIN SODIUM, DICLOXACILLIN SODIUM
 DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
 DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
 DORZOLAMIDE HYDROCHLORIDE AND TIMOLOL MALEATE, DORZOLAMIDE HYDROCHLORIDE
 ENOXAPARIN SODIUM (PRESERVATIVE FREE), ENOXAPARIN SODIUM
 EPLERENONE, EPLERENONE
 ETODOLAC, ETODOLAC
 FLUDARABINE PHOSPHATE, FLUDARABINE PHOSPHATE
 FLUPHENAZINE HYDROCHLORIDE, FLUPHENAZINE HYDROCHLORIDE
 FOSINOPRIL SODIUM AND HYDROCHLOROTHIAZIDE, FOSINOPRIL SODIUM
 FUROSEMIDE, FUROSEMIDE
 GALANTAMINE HYDROBROMIDE, GALANTAMINE HYDROBROMIDE
 GLIPIZIDE, GLIPIZIDE
 HALOPERIDOL, HALOPERIDOL
 HEPARIN SODIUM, HEPARIN SODIUM
 HYDROXYCHLOROQUINE SULFATE, HYDROXYCHLOROQUINE SULFATE
 HYDROXYZINE PAMOATE, HYDROXYZINE PAMOATE
 IMIPRAMINE HYDROCHLORIDE, IMIPRAMINE HYDROCHLORIDE
 INDOMETHACIN, INDOMETHACIN
 IRBESARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 IRBESARTAN, IRBESARTAN
 ISOSORBIDE DINITRATE, ISOSORBIDE DINITRATE
 ITRACONAZOLE, ITRACONAZOLE
 LABETALOL HYDROCHLORIDE, LABETALOL HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

**** S ****

* SANDOZ INC
 LANSOPRAZOLE, LANSOPRAZOLE
 LEUPROLIDE ACETATE, LEUPROLIDE ACETATE
 LEVOFLOXACIN, LEVOFLOXACIN
 LISINAPRIL AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 LORAZEPAM, LORAZEPAM
 LOSARTAN POTASSIUM AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 LOSARTAN POTASSIUM, LOSARTAN POTASSIUM
 LOVASTATIN, LOVASTATIN
 MECLIZINE HYDROCHLORIDE, MECLIZINE HYDROCHLORIDE
 METAXALONE, METAXALONE
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 METHAZOLAMIDE, METHAZOLAMIDE
 METHYLPREDNISOLONE, METHYLPREDNISOLONE
 METOLAZONE, METOLAZONE
 MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE
 MYCOPHENOLATE MOFETIL, MYCOPHENOLATE MOFETIL
 NADOLOL, NADOLOL
 NAFCILLIN SODIUM, NAFCILLIN SODIUM
 NITROFURANTOIN (MONOHYDRATE/MACROCRYSTALS), NITROFURANTOIN
 NIZATIDINE, NIZATIDINE
 OLANZAPINE AND FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 OMEPRAZOLE, OMEPRAZOLE
 OMNITROPE, SOMATROPIN
 ONDANSETRON, ONDANSETRON
 ORPHENADRINE CITRATE, ORPHENADRINE CITRATE
 OXALIPLATIN, OXALIPLATIN
 OXAPROZIN, OXAPROZIN
 OXAZEPAM, OXAZEPAM
 PENICILLIN G POTASSIUM, PENICILLIN G POTASSIUM
 PENICILLIN G SODIUM, PENICILLIN G SODIUM
 PENICILLIN V POTASSIUM, PENICILLIN V POTASSIUM
 PERPHENAZINE, PERPHENAZINE
 PHENTERMINE HYDROCHLORIDE, PHENTERMINE HYDROCHLORIDE
 PIOGLITAZONE HYDROCHLORIDE AND GLIMEPIRIDE, GLIMEPIRIDE
 PIOGLITAZONE HYDROCHLORIDE AND METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 PIOGLITAZONE HYDROCHLORIDE, PIOGLITAZONE HYDROCHLORIDE
 PIPERACILLIN AND TAZOBACTAM, PIPERACILLIN SODIUM
 PROCHLORPERAZINE MALEATE, PROCHLORPERAZINE MALEATE
 PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
 QUETIAPINE FUMARATE, QUETIAPINE FUMARATE
 QUINIDINE SULFATE, QUINIDINE SULFATE
 RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
 RIBAVIRIN, RIBAVIRIN
 RIFAMPIN, RIFAMPIN
 RISPERIDONE, RISPERIDONE
 RIZATRIPTAN BENZOATE, RIZATRIPTAN BENZOATE
 SULFADIAZINE, SULFADIAZINE
 TACROLIMUS, TACROLIMUS
 TAMSULOSIN HYDROCHLORIDE, TAMSULOSIN HYDROCHLORIDE
 TEMAZEPAM, TEMAZEPAM
 TRIAMTERENE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 TRIFLUOPERAZINE HYDROCHLORIDE, TRIFLUOPERAZINE HYDROCHLORIDE
 VALACYCLOVIR HYDROCHLORIDE, VALACYCLOVIR HYDROCHLORIDE
 VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE
 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
 ACETAMINOPHEN, ACETAMINOPHEN
 AMOXICILLIN AND CLAVULANATE POTASSIUM, AMOXICILLIN

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** S **

* SANDOZ INC
 ANECTINE, SUCCINYLCHOLINE CHLORIDE
 ANGIOMAX, BIVALIRUDIN
 ARISTOSPAN, TRIAMCINOLONE HEXACETONIDE
 ASPIRIN AND DIPYRIDAMOLE, ASPIRIN
 ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM
 AZELASTINE HYDROCHLORIDE, AZELASTINE HYDROCHLORIDE
 BETOPTIC, BETAXOLOL HYDROCHLORIDE
 BIMATOPROST, BIMATOPROST
 BRIMONIDINE TARTRATE, BRIMONIDINE TARTRATE
 BROMOCRIPTINE MESYLATE, BROMOCRIPTINE MESYLATE
 BUDESONIDE, BUDESONIDE
 BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
 CARBOPLATIN, CARBOPLATIN
 CARTEOLOL HYDROCHLORIDE, CARTEOLOL HYDROCHLORIDE
 CEFTRIAZONE, CEFTRIAZONE SODIUM
 CISATRACURIUM BESYLATE PRESERVATIVE FREE, CISATRACURIUM BESYLATE
 CISATRACURIUM BESYLATE, CISATRACURIUM BESYLATE
 CLINDAMYCIN PHOSPHATE IN 5% DEXTROSE IN PLASTIC CONTAINER, CLINDAMYCIN PHOSPHATE
 CROMOLYN SODIUM, CROMOLYN SODIUM
 DECITABINE, DECITABINE
 DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE
 DICLOFENAC SODIUM, DICLOFENAC SODIUM
 DIGOXIN, DIGOXIN
 DORZOLAMIDE HYDROCHLORIDE AND TIMOLOL MALEATE, DORZOLAMIDE HYDROCHLORIDE
 DORZOLAMIDE HYDROCHLORIDE, DORZOLAMIDE HYDROCHLORIDE
 DOXERCALCIFEROL, DOXERCALCIFEROL
 ENALAPRIL MALEATE, ENALAPRIL MALEATE
 ENOXAPARIN SODIUM, ENOXAPARIN SODIUM
 EPHEDRINE SULFATE, EPHEDRINE SULFATE
 EZETIMIBE, EZETIMIBE
 FENOLDOPAM MESYLATE, FENOLDOPAM MESYLATE
 FLUMAZENIL, FLUMAZENIL
 FULVESTRANT, FULVESTRANT
 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
 ISONIAZID, ISONIAZID
 KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
 LANSOPRAZOLE, AMOXICILLIN AND CLARITHROMYCIN, AMOXICILLIN
 LATANOPROST, LATANOPROST
 LEVOLEUCOVORIN CALCIUM, LEVOLEUCOVORIN CALCIUM
 LINEZOLID, LINEZOLID
 MAXITROL, DEXAMETHASONE
 MESALAMINE, MESALAMINE
 METHOTREXATE SODIUM PRESERVATIVE FREE, METHOTREXATE SODIUM
 METHYLPREDNISOLONE ACETATE, METHYLPREDNISOLONE ACETATE
 METOPROLOL TARTRATE, METOPROLOL TARTRATE
 MONTELUKAST SODIUM, MONTELUKAST SODIUM
 MYDRIACYL, TROPICAMIDE
 NEOMYCIN AND POLYMYXIN B SULFATES AND HYDROCORTISONE, HYDROCORTISONE
 NEVIRAPINE, NEVIRAPINE
 NOREPINEPHRINE BITARTRATE, NOREPINEPHRINE BITARTRATE
 OLANZAPINE, OLANZAPINE
 ONDANSETRON HYDROCHLORIDE PRESERVATIVE FREE, ONDANSETRON HYDROCHLORIDE
 ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
 PALONOSETRON HYDROCHLORIDE, PALONOSETRON HYDROCHLORIDE
 PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
 PARICALCITOL, PARICALCITOL
 PHENYLEPHRINE HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** S ******* SANDOZ INC**

PIPERACILLIN AND TAZOBACTAM, PIPERACILLIN SODIUM
 PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE
 QOLIANA, BRIMONIDINE TARTRATE
 RASAGILINE MESYLATE, RASAGILINE MESYLATE
 REGONOL, PYRIDOSTIGMINE BROMIDE
 ROCURONIUM BROMIDE, ROCURONIUM BROMIDE
 ROPINIROLE HYDROCHLORIDE, ROPINIROLE HYDROCHLORIDE
 ROSUVASTATIN CALCIUM, ROSUVASTATIN CALCIUM
 SILODOSIN, SILODOSIN
 SULFACETAMIDE SODIUM, SULFACETAMIDE SODIUM
 TELMISARTAN, TELMISARTAN
 TERIFLUNOMIDE, TERIFLUNOMIDE
 TESTOSTERONE CYPIONATE, TESTOSTERONE CYPIONATE
 TIGECYCLINE, TIGECYCLINE
 TIMOLOL MALEATE, TIMOLOL MALEATE
 TOBREX, TOBRAMYCIN
 TREPROSTINIL, TREPROSTINIL
 TRIFLURIDINE, TRIFLURIDINE
 TRIMETHOPRIM SULFATE AND POLYMYXIN B SULFATE, POLYMYXIN B SULFATE
 VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE
 VORICONAZOLE, VORICONAZOLE
 ZIPRASIDONE HYDROCHLORIDE, ZIPRASIDONE HYDROCHLORIDE
 ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE

SANOCHEMIA CORP USA*** SANOCHEMIA CORP USA**

SCANLUX-300, IOPAMIDOL
 SCANLUX-370, IOPAMIDOL

SANOFI*** SANOFI GENZYME**

HECTOROL, DOXERCALCIFEROL
 RENVELA, SEVELAMER CARBONATE

SANOFI AVENTIS US*** SANOFI AVENTIS US INC**

JEVTANA KIT, CABAZITAXEL

*** SANOFI AVENTIS US LLC**

ALLEGRA ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
 ALLEGRA-D 12 HOUR ALLERGY AND CONGESTION, FEXOFENADINE HYDROCHLORIDE (OTC)
 ALLEGRA-D 24 HOUR ALLERGY AND CONGESTION, FEXOFENADINE HYDROCHLORIDE (OTC)
 AMARYL, GLIMEPIRIDE
 AMBIEN CR, ZOLPIDEM TARTRATE
 AMBIEN, ZOLPIDEM TARTRATE
 APIDRA SOLOSTAR, INSULIN GLULISINE RECOMBINANT
 APIDRA, INSULIN GLULISINE RECOMBINANT
 ARAVA, LEFLUNOMIDE
 AUBAGIO, TERIFLUNOMIDE
 AVALIDE, HYDROCHLOROTHIAZIDE
 AVAPRO, IRBESARTAN
 CHILDREN'S ALLEGRA ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
 DIABETA, GLYBURIDE
 ELOXATIN, OXALIPLATIN
 FERRLECIT, SODIUM FERRIC GLUCONATE COMPLEX
 FLOMAX, TAMSULOSIN HYDROCHLORIDE
 GAVISCON, ALUMINUM HYDROXIDE (OTC)
 LANTUS SOLOSTAR, INSULIN GLARGINE RECOMBINANT
 LANTUS, INSULIN GLARGINE RECOMBINANT
 LOVENOX (PRESERVATIVE FREE), ENOXAPARIN SODIUM
 LOVENOX, ENOXAPARIN SODIUM
 MULTAQ, DRONEDARONE HYDROCHLORIDE
 NASACORT ALLERGY 24 HOUR, TRIAMCINOLONE ACETONIDE (OTC)
 NICODERM CQ, NICOTINE (OTC)
 PLAVIX, CLOPIDOGREL BISULFATE
 PRIFTIN, RIFAPENTINE
 PRIMAQUINE, PRIMAQUINE PHOSPHATE
 RIFADIN, RIFAMPIN

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** S ****

* SANOFI AVENTIS US LLC
 RIFAMATE, ISONIAZID
 RIFATER, ISONIAZID
 TAXOTERE, DOCETAXEL
 XYZAL ALLERGY 24HR, LEVOCETIRIZINE DIHYDROCHLORIDE (OTC)
 XYZAL, LEVOCETIRIZINE DIHYDROCHLORIDE

SANOFI US

* SANOFI US
 ZANTAC 150, RANITIDINE HYDROCHLORIDE (OTC)
 ZANTAC 75, RANITIDINE HYDROCHLORIDE (OTC)

SANOFI US SERVICES

* SANOFI US SERVICES INC
 TOUJEO MAX SOLOSTAR, INSULIN GLARGINE RECOMBINANT
 TOUJEO SOLOSTAR, INSULIN GLARGINE RECOMBINANT

SANOFI-AVENTIS US

* SANOFI-AVENTIS US LLC
 ADLYXIN, LIXISENATIDE
 ADMELOG SOLOSTAR, INSULIN LISPRO
 ADMELOG, INSULIN LISPRO
 SOLIQUA 100/33, INSULIN GLARGINE

SANOVEL ILAC

* SANOVEL ILAC SAN VE TIC AS
 ROCURONIUM BROMIDE, ROCURONIUM BROMIDE

SANTARUS INC

* SANTARUS INC
 GLUMETZA, METFORMIN HYDROCHLORIDE

SAOL THERAPS RES LTD

* SAOL THERAPEUTICS RESEARCH LTD
 LIORESAL, BACLOFEN

SAPTALIS PHARMS

* SAPTALIS PHARMACEUTICALS LLC
 CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE

SAREPTA THERAPS INC

* SAREPTA THERAPEUTICS INC
 EXONDYS 51, ETEPLIRSEN
 VYONDYS 53, GOLODIRSEN

SAVIOR LIFETEC CORP

* SAVIOR LIFETEC CORP
 ERTAPENEM SODIUM, ERTAPENEM SODIUM
 MEROPENEM, MEROPENEM

SAWAI USA

* SAWAI USA INC
 PITAVASTATIN CALCIUM, PITAVASTATIN CALCIUM

SB PHARMCO

* SB PHARMCO PUERTO RICO INC
 AVANDIA, ROSIGLITAZONE MALEATE

SCHERING

* SCHERING CORP
 INTEGRILIN, EPTIFIBATIDE
 NOXAFIL, POSACONAZOLE

SCIECURE PHARMA INC

* SCIECURE PHARMA INC
 BUDESONIDE, BUDESONIDE
 PAROXETINE HYDROCHLORIDE, PAROXETINE HYDROCHLORIDE

SCIEGEN PHARMS INC

* SCIEGEN PHARMACEUTICALS INC
 AMLODIPINE AND OLMESARTAN MEDOXOMIL, AMLODIPINE BESYLATE
 ARIPIPIRAZOLE, ARIPIPIRAZOLE
 ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM
 BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
 CARISOPRODOL, CARISOPRODOL
 CLOPIDOGREL BISULFATE, CLOPIDOGREL BISULFATE
 DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** S ******* SCIEGEN PHARMACEUTICALS INC**

ETHACRYNIC ACID, ETHACRYNIC ACID
 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
 LAMOTRIGINE, LAMOTRIGINE
 LEVOCETIRIZINE DIHYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE
 METAXALONE, METAXALONE
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 NABUMETONE, NABUMETONE
 NAPROXEN SODIUM, NAPROXEN SODIUM
 NAPROXEN, NAPROXEN
 OLMESARTAN MEDOXOMIL, OLMESARTAN MEDOXOMIL
 OMEPRAZOLE AND SODIUM BICARBONATE, OMEPRAZOLE
 PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE
 PREGABALIN, PREGABALIN
 QUETIAPINE FUMARATE, QUETIAPINE FUMARATE
 RALOXIFENE HYDROCHLORIDE, RALOXIFENE HYDROCHLORIDE
 RANOLAZINE, RANOLAZINE
 SOLIFENACIN SUCCINATE, SOLIFENACIN SUCCINATE

SCILEX PHARMS INC

*** SCILEX PHARMACEUTICALS**
 ZTLIDO, LIDOCAINE

SCINOPHARM TAIWAN

*** SCINOPHARM TAIWAN LTD**
 FONDAPARINUX SODIUM, FONDAPARINUX SODIUM

SEBELA IRELAND LTD

*** SEBELA IRELAND LTD**
 BRISDELLE, PAROXETINE MESYLATE
 IMURAN, AZATHIOPRINE
 LOTRONEX, ALOSETRON HYDROCHLORIDE
 MICORT-HC, HYDROCORTISONE ACETATE
 MOTOFEN, ATROPINE SULFATE
 NAFTIN, NAFTIFINE HYDROCHLORIDE
 ONMEL, ITRACONAZOLE
 PEXEVA, PAROXETINE MESYLATE
 PRAMOSONE, HYDROCORTISONE ACETATE
 RIDAURA, AURANOFIN

SECAN PHARMS

*** SECAN PHARMACEUTICALS INC**
 LEVETIRACETAM, LEVETIRACETAM

SECURA

*** SECURA BIO INC**
 FARYDAK, PANOBINOSTAT LACTATE

SENTYNL THERAPS INC

*** SENTYNL THERAPEUTICS INC**
 LEVORPHANOL TARTRATE, LEVORPHANOL TARTRATE

SEPTODONT

*** SEPTODONT INC**
 BUPIVACAINE HYDROCHLORIDE AND EPINEPHRINE, BUPIVACAINE HYDROCHLORIDE

SEPTODONT HOLDING

*** SEPTODONT HOLDING SAS**
 ORAVERSE, PHENTOLAMINE MESYLATE

SEPTODONT INC

*** SEPTODONT INC**
 LIDOCAINE, LIDOCAINE
 PRILOCAINE HYDROCHLORIDE AND EPINEPHRINE BITARTRATE, EPINEPHRINE BITARTRATE
 PRILOCAINE HYDROCHLORIDE, PRILOCAINE HYDROCHLORIDE

SERB SA

*** SERB SA**

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** S ****

* SERB SA

CYANOKIT, HYDROXOCOBALAMIN

SETON PHARM

* SETON PHARMACEUTICAL LLC

PEDIAPRED, PREDNISOLONE SODIUM PHOSPHATE

SETON PHARMS

* SETON PHARMACEUTICALS LLC

CAPTOPRIL, CAPTOPRIL

PENTAMIDINE ISETHIONATE, PENTAMIDINE ISETHIONATE

SHANDONG

* SHANDONG NEW TIME PHARMACEUTICAL CO LTD

ROSUVASTATIN CALCIUM, ROSUVASTATIN CALCIUM

SHANDONG XINHUA

* SHANDONG XINHUA PHARMACEUTICAL CO LTD

IBUPROFEN, IBUPROFEN

IBUPROFEN, IBUPROFEN (OTC)

SHANGHAI HENGRUI

* SHANGHAI HENGRUI PHARMACEUTICAL CO LTD

DESFLURANE, DESFLURANE

SEVOFLURANE, SEVOFLURANE

SHENZHEN TECHDOW

* SHENZHEN TECHDOW PHARMACEUTICAL CO LTD

HEPARIN SODIUM PRESERVATIVE FREE, HEPARIN SODIUM

HEPARIN SODIUM, HEPARIN SODIUM

SHERTECH LABS LLC

* SHERTECH LABORATORIES LLC

AMMONIA N 13, AMMONIA N-13

FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18

SODIUM FLUORIDE F-18, SODIUM FLUORIDE F-18

SHIELD TX

* SHIELD TX UK LTD

ACCRUFER, FERRIC MALTOL

SHILPA MEDICARE

* SHILPA MEDICARE LTD

AZACITIDINE, AZACITIDINE

SHILPA MEDICARE LTD

* SHILPA MEDICARE LTD

BUSULFAN, BUSULFAN

CAPECITABINE, CAPECITABINE

DOCETAXEL, DOCETAXEL

ERLOTINIB HYDROCHLORIDE, ERLOTINIB HYDROCHLORIDE

GEMCITABINE HYDROCHLORIDE, GEMCITABINE HYDROCHLORIDE

IMATINIB MESYLATE, IMATINIB MESYLATE

IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE

SHIONOGI INC

* SHIONOGI INC

FETROJA, CEFIDEROCOL SULFATE TOSYLATE

MULPLETA, LUSUTROMBOPAG

PONSTEL, MEFENAMIC ACID

SHIRE

* SHIRE DEVELOPMENT INC

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

INTUNIV, GUANFACINE HYDROCHLORIDE

LIALDA, MESALAMINE

PENTASA, MESALAMINE

SHIRE DEV LLC

* SHIRE DEVELOPMENT LLC

CARBATROL, CARBAMAZEPINE

FOSRENOL, LANTHANUM CARBONATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** S ****

* SHIRE DEVELOPMENT LLC
 MOTEGRITY, PRUCALOPRIDE SUCCINATE
 MYDAYIS, AMPHETAMINE ASPARTATE
 VYVANSE, LISDEXAMFETAMINE DIMESYLATE

SHIRE DEVELOPMENT

* SHIRE DEVELOPMENT INC
 VYVANSE, LISDEXAMFETAMINE DIMESYLATE

SHIRE HUMAN GENETIC

* SHIRE HUMAN GENETIC THERAPIES INC
 VPRIV, VELAGLUCERASE ALFA

SHIRE LLC

* SHIRE DEVELOPMENT LLC
 AGRYLIN, ANAGRELIDE HYDROCHLORIDE
 FOSRENOL, LANTHANUM CARBONATE

SHIRE ORPHAN THERAP

* SHIRE ORPHAN THERAPIES LLC
 FIRAZYR, ICATIBANT ACETATE

SIDMAK LABS INDIA

* SIDMAK LABORATORIES INDIA PVT LTD
 MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE

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
 DISULFIRAM, DISULFIRAM
 DOFETILIDE, DOFETILIDE
 ERGOCALCIFEROL, ERGOCALCIFEROL
 FLUCYTOSINE, FLUCYTOSINE
 GRISEOFULVIN, GRISEOFULVIN, MICROSIZE
 GRISEOFULVIN, ULTRAMICROSIZE, GRISEOFULVIN, ULTRAMICROSIZE
 LIOTHYRONINE SODIUM, LIOTHYRONINE SODIUM
 NITROGLYCERIN, NITROGLYCERIN
 POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
 PROTRIPTYLINE HYDROCHLORIDE, PROTRIPTYLINE HYDROCHLORIDE
 SODIUM PHENYL BUTYRATE, SODIUM PHENYL BUTYRATE

SILVERGATE PHARMS

* SILVERGATE PHARMACEUTICALS INC
 EPANED, ENALAPRIL MALEATE
 KATERZIA, AMLODIPINE BENZOATE
 QBRELIS, LISINAPRIL
 XATMEP, METHOTREXATE SODIUM

SINOTHERAPEUTICS INC

* SINOTHERAPEUTICS INC
 BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
 POSACONAZOLE, POSACONAZOLE
 PROPAFENONE HYDROCHLORIDE, PROPAFENONE HYDROCHLORIDE

SKINMEDICA

* SKINMEDICA INC
 VANIQA, EFLORNITHINE HYDROCHLORIDE

SKYEPHARMA AG

* SKYEPHARMA AG
 TRIGLIDE, FENOFIBRATE

SLATE

* SLATE RUN PHARMACEUTICALS LLC
 CILOSTAZOL, CILOSTAZOL
 DICYCLOMINE HYDROCHLORIDE, DICYCLOMINE HYDROCHLORIDE
 METHOCARBAMOL, METHOCARBAMOL

SLAYBACK PHARMA LLC

* SLAYBACK PHARMA LLC

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** S ****

- * SLAYBACK PHARMA LLC
DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE
HYDROXYPROGESTERONE CAPROATE, HYDROXYPROGESTERONE CAPROATE

SMITHKLINE BEECHAM

- * SMITHKLINE BEECHAM
LOVAZA, OMEGA-3-ACID ETHYL ESTERS
- * SMITHKLINE BEECHAM (CORK) LTD IRELAND
COREG CR, CARVEDILOL PHOSPHATE
COREG, CARVEDILOL

SOAPCO

- * SOAPCO INC
BRIAN CARE, CHLORHEXIDINE GLUCONATE (OTC)

SOFGEN PHARMS

- * SOFGEN PHARMACEUTICALS
NIMODIPINE, NIMODIPINE
- * 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
FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18
SODIUM FLUORIDE F-18, SODIUM FLUORIDE F-18
- * SOFIE CO DBA SOFIE (FKA ZEVACOR PHARMA INC)
AMMONIA N 13, AMMONIA N-13
FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18
SODIUM FLUORIDE F-18, SODIUM FLUORIDE F-18

SOMERSET

- * SOMERSET PHARMACEUTICALS INC
EMSAM, SELEGILINE
- * SOMERSET THERAPEUTICS LLC
CISATRACURIUM BESYLATE, CISATRACURIUM BESYLATE
DEXAMETHASONE SODIUM PHOSPHATE, DEXAMETHASONE SODIUM PHOSPHATE
DORZOLAMIDE HYDROCHLORIDE AND TIMOLOL MALEATE, DORZOLAMIDE HYDROCHLORIDE
LATANOPROST, LATANOPROST
SULFAMETHOXAZOLE AND TRIMETHOPRIM, SULFAMETHOXAZOLE
VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE

SOMERSET THERAPS LLC

- * SOMERSET THERAPEUTICS LLC
AZELASTINE HYDROCHLORIDE, AZELASTINE HYDROCHLORIDE
BIMATOPROST, BIMATOPROST
BRIMONIDINE TARTRATE, BRIMONIDINE TARTRATE
CISATRACURIUM BESYLATE PRESERVATIVE FREE, CISATRACURIUM BESYLATE
CYANOCOBALAMIN, CYANOCOBALAMIN
DEXAMETHASONE SODIUM PHOSPHATE PRESERVATIVE FREE, DEXAMETHASONE SODIUM PHOSPHATE
DEXAMETHASONE SODIUM PHOSPHATE, DEXAMETHASONE SODIUM PHOSPHATE
EPINASTINE HYDROCHLORIDE, EPINASTINE HYDROCHLORIDE
GLYCOPYRROLATE, GLYCOPYRROLATE
HALOPERIDOL DECANOATE, HALOPERIDOL DECANOATE
METHOCARBAMOL, METHOCARBAMOL
NALOXONE HYDROCHLORIDE, NALOXONE HYDROCHLORIDE
OLOPATADINE HYDROCHLORIDE, OLOPATADINE HYDROCHLORIDE
PILOCARPINE HYDROCHLORIDE, PILOCARPINE HYDROCHLORIDE
ROPIVACAINE HYDROCHLORIDE, ROPIVACAINE HYDROCHLORIDE
SODIUM NITROPRUSSIDE, SODIUM NITROPRUSSIDE
SUCCINYLCHOLINE CHLORIDE, SUCCINYLCHOLINE CHLORIDE
TOBRAMYCIN, TOBRAMYCIN
TROPICAMIDE, TROPICAMIDE
VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE

SOMMER PHARMS II LLC

- * SOMMER PHARMACEUTICALS II LLC
TECHNETIUM TC99M MERTIATIDE KIT, TECHNETIUM TC-99M MERTIATIDE KIT

SPARC

- * SUN PHARMA ADVANCED RESEARCH CO LTD

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** S ****

* SUN PHARMA ADVANCED RESEARCH CO LTD
ELEPSIA XR, LEVETIRACETAM

SPECGX LLC

* SPECGX LLC
ACETAMINOPHEN AND CODEINE PHOSPHATE, ACETAMINOPHEN
ANAFRANIL, CLOMIPRAMINE HYDROCHLORIDE
ANEXSIA 5/325, ACETAMINOPHEN
ANEXSIA 7.5/325, ACETAMINOPHEN
BENZPHETAMINE HYDROCHLORIDE, BENZPHETAMINE HYDROCHLORIDE
BUPRENORPHINE HYDROCHLORIDE AND NALOXONE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE
BUTALBITAL, ACETAMINOPHEN AND CAFFEINE, ACETAMINOPHEN
DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE
DEXTROAMPHETAMINE SULFATE, DEXTROAMPHETAMINE SULFATE
FENTANYL CITRATE, FENTANYL CITRATE
FENTANYL-100, FENTANYL
FENTANYL-12, FENTANYL
FENTANYL-25, FENTANYL
FENTANYL-50, FENTANYL
FENTANYL-75, FENTANYL
FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
HYDROMORPHONE HYDROCHLORIDE, HYDROMORPHONE HYDROCHLORIDE
IMIPRAMINE HYDROCHLORIDE, IMIPRAMINE HYDROCHLORIDE
LEVORPHANOL TARTRATE, LEVORPHANOL TARTRATE
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
OXYMORPHONE HYDROCHLORIDE, OXYMORPHONE HYDROCHLORIDE
PAMELOR, NORTRIPTYLINE HYDROCHLORIDE
RESTORIL, TEMAZEPAM
ROXICODONE, OXYCODONE HYDROCHLORIDE
TOFRANIL, IMIPRAMINE HYDROCHLORIDE

SPECTRA MDCL DEVICES

* SPECTRA MEDICAL DEVICES INC
LIDOCAINE HYDROCHLORIDE, LIDOCAINE HYDROCHLORIDE
SODIUM CHLORIDE 0.9%, SODIUM CHLORIDE

SPECTRON MRC LLC

* SPECTRON MRC LLC
AMMONIA N 13, AMMONIA N-13
FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18
SODIUM FLUORIDE F-18, SODIUM FLUORIDE F-18

SPIL

* SUN PHARMA INDUSTRIES LTD
KAPSPARGO SPRINKLE, METOPROLOL SUCCINATE
NIFEDIPINE, NIFEDIPINE
OMEPRAZOLE MAGNESIUM, OMEPRAZOLE MAGNESIUM (OTC)

SPROUT PHARMS

* SPROUT PHARMACEUTICALS INC
ADDYI, FLIBANSERIN

SQUARE PHARMS LTD

* SQUARE PHARMACEUTICALS LTD
ACYCLOVIR, ACYCLOVIR
VALSARTAN, VALSARTAN

ST RENATUS

* ST RENATUS LLC
KOVANAZE, OXYMETAZOLINE HYDROCHLORIDE

STAND HOMEOPATH

* STANDARD HOMEOPATHIC CO

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** S ****

* STANDARD HOMEOPATHIC CO
 IVY BLOCK, BENTOQUATAM (OTC)

STANDARD CHEM PHARM

* STANDARD CHEM AND PHARM CO LTD
 RILUZOLE, RILUZOLE

STASON PHARMS

* STASON PHARMACEUTICALS INC
 PURINETHOL, MERCAPTOPYRINE

STI PHARMA LLC

* STI PHARMA LLC
 ARSENIC TRIOXIDE, ARSENIC TRIOXIDE
 CARMUSTINE, CARMUSTINE
 CYCLOPHOSPHAMIDE, CYCLOPHOSPHAMIDE
 MYAMBUTOL, ETHAMBUTOL HYDROCHLORIDE
 THIOTEPA, THIOTEPA
 TRIACIN-C, CODEINE PHOSPHATE

STIEFEL

* STIEFEL LABORATORIES INC
 DUAC, BENZOYL PEROXIDE

STIEFEL LABS INC

* STIEFEL LABORATORIES INC
 SORIATANE, ACITRETIN

STRIDES PHARMA

* STRIDES PHARMA GLOBAL PTE LTD
 ABACAVIR SULFATE, ABACAVIR SULFATE
 ACARBOSE, ACARBOSE
 ACETAZOLAMIDE, ACETAZOLAMIDE
 ALBENDAZOLE, ALBENDAZOLE
 AMANTADINE HYDROCHLORIDE, AMANTADINE HYDROCHLORIDE
 AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
 BENZONATATE, BENZONATATE
 BUSPIRONE HYDROCHLORIDE, BUSPIRONE HYDROCHLORIDE
 CALCITRIOL, CALCITRIOL
 CARISOPRODOL, CARISOPRODOL
 CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 CINACALCET HYDROCHLORIDE, CINACALCET HYDROCHLORIDE
 CLARITHROMYCIN, CLARITHROMYCIN
 CYPROHEPTADINE HYDROCHLORIDE, CYPROHEPTADINE HYDROCHLORIDE
 DICLOFENAC POTASSIUM, DICLOFENAC POTASSIUM
 DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
 DUTASTERIDE, DUTASTERIDE
 EFAVIRENZ, EFAVIRENZ
 ERGOCALCIFEROL, ERGOCALCIFEROL
 ETHOSUXIMIDE, ETHOSUXIMIDE
 GABAPENTIN, GABAPENTIN
 HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
 HYDROCORTISONE, HYDROCORTISONE
 IBUPROFEN AND DIPHENHYDRAMINE HYDROCHLORIDE, DIPHENHYDRAMINE HYDROCHLORIDE (OTC)
 IBUPROFEN, IBUPROFEN
 IBUPROFEN, IBUPROFEN (OTC)
 KETOCONAZOLE, KETOCONAZOLE
 LAMIVUDINE AND ZIDOVUDINE, LAMIVUDINE
 LAMIVUDINE, LAMIVUDINE
 LIDOCAINE, LIDOCAINE
 LORATADINE, LORATADINE (OTC)
 LOSARTAN POTASSIUM, LOSARTAN POTASSIUM
 MELOXICAM, MELOXICAM
 MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
 METHOXSALEN, METHOXSALEN
 METRONIDAZOLE, METRONIDAZOLE
 MYCOPHENOLATE MOFETIL, MYCOPHENOLATE MOFETIL
 NEVIRAPINE, NEVIRAPINE
 NYSTATIN AND TRIAMCINOLONE ACETONIDE, NYSTATIN
 OMEGA-3-ACID ETHYL ESTERS, OMEGA-3-ACID ETHYL ESTERS
 OSELTAMIVIR PHOSPHATE, OSELTAMIVIR PHOSPHATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** S ****

* STRIDES PHARMA GLOBAL PTE LTD
 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
 PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
 RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
 RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE (OTC)
 SOLIFENACIN SUCCINATE, SOLIFENACIN SUCCINATE
 TACROLIMUS, TACROLIMUS
 TENOFOVIR DISOPROXIL FUMARATE, TENOFOVIR DISOPROXIL FUMARATE
 TETRACYCLINE HYDROCHLORIDE, TETRACYCLINE HYDROCHLORIDE
 TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE
 VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE

STRONGBRIDGE US

* STRONGBRIDGE US INC
 KEVEYIS, DICHLORPHENAMIDE

SUCAMPO PHARMA LLC

* SUCAMPO PHARMA AMERICAS LLC
 AMITIZA, LUBIPROSTONE

SUN PHARM

* SUN PHARMACEUTICAL INDUSTRIES LTD
 ABSORICA LD, ISOTRETINOIN
 ALBUTEROL SULFATE AND IPRATROPIUM BROMIDE, ALBUTEROL SULFATE
 ALBUTEROL SULFATE, ALBUTEROL SULFATE
 ALENDRONATE SODIUM, ALENDRONATE SODIUM
 ALFUZOSIN HYDROCHLORIDE, ALFUZOSIN HYDROCHLORIDE
 ALPRAZOLAM, ALPRAZOLAM
 AMBRISENTAN, AMBRISENTAN
 AMIFOSTINE, AMIFOSTINE
 ASPIRIN AND DIPYRIDAMOLE, ASPIRIN
 AZELASTINE HYDROCHLORIDE, AZELASTINE HYDROCHLORIDE
 BICALUTAMIDE, BICALUTAMIDE
 BOSENTAN, BOSENTAN
 BUPRENORPHINE HYDROCHLORIDE AND NALOXONE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE
 BUPRENORPHINE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE
 BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
 BYNFEZIA PEN, OCTREOTIDE ACETATE
 CAFFEINE CITRATE, CAFFEINE CITRATE
 CAPECITABINE, CAPECITABINE
 CARBIDOPA AND LEVODOPA, CARBIDOPA
 CARBIDOPA, LEVODOPA AND ENTACAPONE, CARBIDOPA
 CARBOPLATIN, CARBOPLATIN
 CERINTA, ETHINYL ESTRADIOL
 CHILDREN'S CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 CHILDREN'S CETIRIZINE HYDROCHLORIDE HIVES RELIEF, CETIRIZINE HYDROCHLORIDE (OTC)
 CHLOROTHIAZIDE SODIUM, CHLOROTHIAZIDE SODIUM
 CINACALCET HYDROCHLORIDE, CINACALCET HYDROCHLORIDE
 CLOPIDOGREL BISULFATE, CLOPIDOGREL BISULFATE
 DALFAMPRIDINE, DALFAMPRIDINE
 DEFERASIROX, DEFERASIROX
 DESMOPRESSIN ACETATE (NEEDS NO REFRIGERATION), DESMOPRESSIN ACETATE
 DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
 DOFETILIDE, DOFETILIDE
 DOXERCALCIFEROL, DOXERCALCIFEROL
 DOXORUBICIN HYDROCHLORIDE (LIPOSOMAL), DOXORUBICIN HYDROCHLORIDE
 DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE
 ENTACAPONE, ENTACAPONE
 ERLOTINIB HYDROCHLORIDE, ERLOTINIB HYDROCHLORIDE
 ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM
 ESZOPICLONE, ESZOPICLONE
 FEBUXOSTAT, FEBUXOSTAT

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** S ****

* SUN PHARMACEUTICAL INDUSTRIES LTD
 FENOFIBRATE, FENOFIBRATE
 FEXOFENADINE HYDROCHLORIDE AND PSEUDOEPHEDRINE HYDROCHLORIDE, FEXOFENADINE
 FINASTERIDE, FINASTERIDE
 FINGOLIMOD HYDROCHLORIDE, FINGOLIMOD HYDROCHLORIDE
 FOSAMPRENAVIR CALCIUM, FOSAMPRENAVIR CALCIUM
 FOSPHENYTOIN SODIUM, FOSPHENYTOIN SODIUM
 GALANTAMINE HYDROBROMIDE, GALANTAMINE HYDROBROMIDE
 GANIRELIX ACETATE, GANIRELIX ACETATE
 GEMCITABINE HYDROCHLORIDE, GEMCITABINE HYDROCHLORIDE
 GUANFACINE HYDROCHLORIDE, GUANFACINE HYDROCHLORIDE
 HYDROXYPROGESTERONE CAPROATE, HYDROXYPROGESTERONE CAPROATE
 IBANDRONATE SODIUM, IBANDRONATE SODIUM
 IMATINIB MESYLATE, IMATINIB MESYLATE
 INFUGEM, GEMCITABINE HYDROCHLORIDE
 IPRATROPIUM BROMIDE, IPRATROPIUM BROMIDE
 KEMEYA, DROSPIRENONE
 KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
 KYRA, DROSPIRENONE
 LANSOPRAZOLE, LANSOPRAZOLE
 LEUPROLIDE ACETATE, LEUPROLIDE ACETATE
 LEVALBUTEROL HYDROCHLORIDE, LEVALBUTEROL HYDROCHLORIDE
 LEVETIRACETAM, LEVETIRACETAM
 LEVOCETIRIZINE DIHYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE
 LIOTHYRONINE SODIUM, LIOTHYRONINE SODIUM
 LORATADINE, LORATADINE (OTC)
 LURASIDONE HYDROCHLORIDE, LURASIDONE HYDROCHLORIDE
 MEDROXYPROGESTERONE ACETATE, MEDROXYPROGESTERONE ACETATE
 MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
 MESALAMINE, MESALAMINE
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 METHOTREXATE SODIUM, METHOTREXATE SODIUM
 NALTREXONE HYDROCHLORIDE, NALTREXONE HYDROCHLORIDE
 NIACIN, NIACIN
 OMEGA-3-ACID ETHYL ESTERS, OMEGA-3-ACID ETHYL ESTERS
 OMEPRAZOLE, OMEPRAZOLE (OTC)
 OPCICON ONE-STEP, LEVONORGESTREL (OTC)
 PALIPERIDONE, PALIPERIDONE
 PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
 PREGABALIN, PREGABALIN
 QUETIAPINE FUMARATE, QUETIAPINE FUMARATE
 RANOLAZINE, RANOLAZINE
 RIOMET ER, METFORMIN HYDROCHLORIDE
 RISEDRONATE SODIUM, RISEDRONATE SODIUM
 RIVASTIGMINE TARTRATE, RIVASTIGMINE TARTRATE
 ROSUVASTATIN CALCIUM, ROSUVASTATIN CALCIUM
 SUMATRIPTAN AND NAPROXEN SODIUM, NAPROXEN SODIUM
 SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
 SYLEVIA, ETHINYL ESTRADIOL
 TADALAFIL, TADALAFIL
 TEMOZOLOMIDE, TEMOZOLOMIDE
 TETRABENAZINE, TETRABENAZINE
 TOBRAMYCIN, TOBRAMYCIN
 TOPIRAMATE, TOPIRAMATE
 TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE
 VECURONIUM BROMIDE, VECURONIUM BROMIDE
 VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE
 ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE

SUN PHARM INDS

* SUN PHARMACEUTICAL INDUSTRIES LTD
 CARBIDOPA AND LEVODOPA, CARBIDOPA
 CHILDREN'S FEXOFENADINE HYDROCHLORIDE ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
 CHILDREN'S FEXOFENADINE HYDROCHLORIDE HIVES, FEXOFENADINE HYDROCHLORIDE (OTC)
 DESLORATADINE, DESLORATADINE
 DIVALPROEX SODIUM, DIVALPROEX SODIUM

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** S ****

* SUN PHARMACEUTICAL INDUSTRIES LTD
 DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
 DOXORUBICIN HYDROCHLORIDE, DOXORUBICIN HYDROCHLORIDE
 EXTENDED PHENYTOIN SODIUM, PHENYTOIN SODIUM
 FEXOFENADINE HYDROCHLORIDE ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
 FEXOFENADINE HYDROCHLORIDE HIVES, FEXOFENADINE HYDROCHLORIDE (OTC)
 METOPROLOL TARTRATE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 OLANZAPINE, OLANZAPINE
 ONDANSETRON, ONDANSETRON
 OXCARBAZEPINE, OXCARBAZEPINE
 SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
 TIAGABINE HYDROCHLORIDE, TIAGABINE HYDROCHLORIDE

SUN PHARM INDS (IN)

* SUN PHARMACEUTICAL INDUSTRIES LTD
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
 ZONISAMIDE, ZONISAMIDE

SUN PHARM INDS INC

* SUN PHARMACEUTICAL INDUSTRIES INC
 ABSORICA, ISOTRETINOIN
 AMITRIPTYLINE HYDROCHLORIDE, AMITRIPTYLINE HYDROCHLORIDE
 CLONAZEPAM, CLONAZEPAM
 CLOZAPINE, CLOZAPINE
 DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE
 DIGOXIN, DIGOXIN
 ERGOCALCIFEROL, ERGOCALCIFEROL
 EURAX, CROTAMITON
 FLUMADINE, RIMANTADINE HYDROCHLORIDE
 GLIPIZIDE, GLIPIZIDE
 HALOG, HALCINONIDE
 HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
 ISOSORBIDE DINITRATE, ISOSORBIDE DINITRATE
 KENALOG, TRIAMCINOLONE ACETONIDE
 LITHIUM CARBONATE, LITHIUM CARBONATE
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
 METOPROLOL TARTRATE, METOPROLOL TARTRATE
 MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE
 MIRTAZAPINE, MIRTAZAPINE
 OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN
 OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
 PAROMOMYCIN SULFATE, PAROMOMYCIN SULFATE
 PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
 REPAGLINIDE, REPAGLINIDE
 TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE
 TRAMADOL HYDROCHLORIDE AND ACETAMINOPHEN, ACETAMINOPHEN
 TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE
 ULTRAVATE, HALOBETASOL PROPIONATE
 VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE
 VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE

SUN PHARM INDS LTD

* SUN PHARMACEUTICAL INDUSTRIES LTD
 ACETAMINOPHEN AND CODEINE PHOSPHATE, ACETAMINOPHEN
 ACETAMINOPHEN, ACETAMINOPHEN (OTC)
 AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
 ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM
 AZITHROMYCIN, AZITHROMYCIN
 BENAZEPRIL HYDROCHLORIDE, BENAZEPRIL HYDROCHLORIDE
 CARVEDILOL, CARVEDILOL
 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

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** S ****

* SUN PHARMACEUTICAL INDUSTRIES LTD
 DESMOPRESSIN ACETATE, DESMOPRESSIN ACETATE
 DOXYCYCLINE, DOXYCYCLINE
 FAMOTIDINE, FAMOTIDINE (OTC)
 FELODIPINE, FELODIPINE
 FENOFIBRATE, FENOFIBRATE
 FLECAINIDE ACETATE, FLECAINIDE ACETATE
 FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 GABAPENTIN, GABAPENTIN
 GLYCOPYRROLATE, GLYCOPYRROLATE
 LEVETIRACETAM, LEVETIRACETAM
 LEVOCETIRIZINE DIHYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE
 LISINAPRIL AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 LISINAPRIL, LISINAPRIL
 LOPERAMIDE HYDROCHLORIDE AND SIMETHICONE, LOPERAMIDE HYDROCHLORIDE (OTC)
 LORATADINE AND PSEUDOEPHEDRINE SULFATE, LORATADINE (OTC)
 LORATADINE REDIDOSE, LORATADINE (OTC)
 LORATADINE, LORATADINE (OTC)
 MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE
 MORPHINE SULFATE, MORPHINE SULFATE
 NALOXONE HYDROCHLORIDE AND PENTAZOCINE HYDROCHLORIDE, NALOXONE HYDROCHLORIDE
 NAPROXEN SODIUM, NAPROXEN SODIUM (OTC)
 NARATRIPTAN, NARATRIPTAN HYDROCHLORIDE
 ONDANSETRON, ONDANSETRON
 OXCARBAZEPINE, OXCARBAZEPINE
 PSEUDOEPHEDRINE HYDROCHLORIDE, PSEUDOEPHEDRINE HYDROCHLORIDE (OTC)
 RILUZOLE, RILUZOLE
 RISPERIDONE, RISPERIDONE
 SERTRALINE HYDROCHLORIDE, SERTRALINE HYDROCHLORIDE
 SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
 TAMSULOSIN HYDROCHLORIDE, TAMSULOSIN HYDROCHLORIDE
 TESTOSTERONE CYPIONATE, TESTOSTERONE CYPIONATE
 TOPIRAMATE, TOPIRAMATE
 VALACYCLOVIR HYDROCHLORIDE, VALACYCLOVIR HYDROCHLORIDE
 VALPROIC ACID, VALPROIC ACID

SUN PHARM INDUSTRIES

* SUN PHARMACEUTICAL INDUSTRIES INC
 ALBUTEROL SULFATE, ALBUTEROL SULFATE
 ALLOPURINOL, ALLOPURINOL
 BACTRIM DS, SULFAMETHOXAZOLE
 BACTRIM, SULFAMETHOXAZOLE
 CARVEDILOL PHOSPHATE, CARVEDILOL PHOSPHATE
 CHLORTHALIDONE, CHLORTHALIDONE
 DEXMETHYLPHENIDATE HYDROCHLORIDE, DEXMETHYLPHENIDATE HYDROCHLORIDE
 DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE
 DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
 ERGOLOID MESYLATES, ERGOLOID MESYLATES
 FELODIPINE, FELODIPINE
 IMIPRAMINE HYDROCHLORIDE, IMIPRAMINE HYDROCHLORIDE
 METHADONE HYDROCHLORIDE, METHADONE HYDROCHLORIDE
 MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE
 MINOXIDIL, MINOXIDIL
 NITROFURANTOIN, NITROFURANTOIN, MACROCRYSTALLINE
 NYSTATIN, NYSTATIN
 PHENTERMINE HYDROCHLORIDE, PHENTERMINE HYDROCHLORIDE
 PINDOLOL, PINDOLOL
 PREDNISONE, PREDNISONE
 PROPAFENONE HYDROCHLORIDE, PROPAFENONE HYDROCHLORIDE
 QUALAQUIN, QUININE SULFATE
 QUINIDINE GLUCONATE, QUINIDINE GLUCONATE
 SPIRONOLACTONE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 SPIRONOLACTONE, SPIRONOLACTONE
 SULFAMETHOXAZOLE AND TRIMETHOPRIM, SULFAMETHOXAZOLE
 SULINDAC, SULINDAC
 TEMAZEPAM, TEMAZEPAM

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** s ****

* SUN PHARMACEUTICAL INDUSTRIES INC
 TRAZODONE HYDROCHLORIDE, TRAZODONE HYDROCHLORIDE
 TRIMETHOBENZAMIDE HYDROCHLORIDE, TRIMETHOBENZAMIDE HYDROCHLORIDE
 ULTRAVATE, HALOBETASOL PROPIONATE

SUN PHARMA GLOBAL

* SUN PHARMA GLOBAL FZE
 BROMSITE, BROMFENAC SODIUM
 CEQUA, CYCLOSPORINE
 DECITABINE, DECITABINE
 DOCETAXEL, DOCETAXEL
 DRIZALMA SPRINKLE, DULOXETINE HYDROCHLORIDE
 ESOMEPRAZOLE SODIUM, ESOMEPRAZOLE SODIUM
 EZALLOR, ROSUVASTATIN CALCIUM
 ODOMZO, SONIDEGIB PHOSPHATE
 ORTIKOS, BUDESONIDE
 XELPROS, LATANOPROST
 YONSA, ABIRATERONE ACETATE

SUNGEN PHARMA

* SUNGEN PHARMA LLC
 DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE
 DICYCLOMINE HYDROCHLORIDE, DICYCLOMINE HYDROCHLORIDE
 FOSAPREPITANT DIMEGLUMINE, FOSAPREPITANT DIMEGLUMINE
 LIDOCAINE, LIDOCAINE
 METHYLPREDNISOLONE, METHYLPREDNISOLONE

SUNNY PHARMTECH INC

* SUNNY PHARMTECH INC
 AMINOCAPROIC ACID, AMINOCAPROIC ACID
 NITROFURANTOIN (MONOHYDRATE/MACROCRYSTALS), NITROFURANTOIN

SUNOVION

* SUNOVION PHARMACEUTICALS INC
 BROVANA, ARFORMOTEROL TARTRATE
 XOPENEX HFA, LEVALBUTEROL TARTRATE

SUNOVION PHARMS INC

* SUNOVION PHARMACEUTICALS INC
 APTIOM, ESLICARBAZEPINE ACETATE
 ARCAPTA NEOHALER, INDACATEROL MALEATE
 LATUDA, LURASIDONE HYDROCHLORIDE
 LUNESTA, ESZOPICLONE
 SEEBRI, GLYCOPYRROLATE
 UTIBRON, GLYCOPYRROLATE
 ZONEGRAN, ZONISAMIDE

SUNOVION RESP

* SUNOVION RESPIRATORY DEVELOPMENT INC
 LONHALA MAGNAIR KIT, GLYCOPYRROLATE

SUNRISE PHARM INC

* SUNRISE PHARMACEUTICAL INC
 NAPROXEN, NAPROXEN

SUNSHINE LAKE

* SUNSHINE LAKE PHARMA CO LTD
 AZITHROMYCIN, AZITHROMYCIN
 CLARITHROMYCIN, CLARITHROMYCIN
 ENTACAPONE, ENTACAPONE
 OLMESARTAN MEDOXOMIL, OLMESARTAN MEDOXOMIL
 OSELTAMIVIR PHOSPHATE, OSELTAMIVIR PHOSPHATE
 TADALAFIL, TADALAFIL

SUNSTAR AMERICAS

* SUNSTAR AMERICAS INC
 PAROEX, CHLORHEXIDINE GLUCONATE

SUPERNUS PHARMS

* SUPERNUS PHARMACEUTICALS INC
 OXTELLAR XR, OXCARBAZEPINE
 TROKENDI XR, TOPIRAMATE

SUVEN LIFE

* SUVEN LIFE SCIENCES LTD

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** S ****

* SUVEN LIFE SCIENCES LTD
 CALCIUM ACETATE, CALCIUM ACETATE
 MALATHION, MALATHION

SVC PHARMA

* SVC PHARMA LP
 DRONABINOL, DRONABINOL

SWEDISH ORPHAN

* SWEDISH ORPHAN BIOVITRUM AB PUBL
 ORFADIN, NITISINONE

SYNTHON PHARMS

* SYNTHON PHARMACEUTICALS INC
 LEVOCETIRIZINE DIHYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE
 TAMSULOSIN HYDROCHLORIDE, TAMSULOSIN HYDROCHLORIDE

**** 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 HOSP RES

* THE METHODIST HOSP RESEARCH INSTITUTE
 FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18

RITEDOSE CORP

* THE RITEDOSE CORP
 ALBUTEROL SULFATE AND IPRATROPIUM BROMIDE, ALBUTEROL SULFATE
 ALBUTEROL SULFATE, ALBUTEROL SULFATE
 IPRATROPIUM BROMIDE, IPRATROPIUM BROMIDE
 LEVALBUTEROL HYDROCHLORIDE, LEVALBUTEROL HYDROCHLORIDE

TAIHO ONCOLOGY

* TAIHO ONCOLOGY INC
 LONSURF, TIPIRACIL HYDROCHLORIDE

TAKEDA PHARMS USA

* TAKEDA PHARMACEUTICALS USA INC
 ACTOPLUS MET, METFORMIN HYDROCHLORIDE
 ACTOS, PIOGLITAZONE HYDROCHLORIDE
 COLCRYS, COLCHICINE
 DEXILANT, DEXLANSOPRAZOLE
 DUEFACT, GLIMEPIRIDE
 KAZANO, ALOGLIPTIN BENZOATE
 NESINA, ALOGLIPTIN BENZOATE
 OSENI, ALOGLIPTIN BENZOATE
 PREVACID, LANSOPRAZOLE
 ROZEREM, RAMELTEON
 TRINTELLIX, VORTIOXETINE HYDROBROMIDE
 ULORIC, FEBUXOSTAT

TAMARANG

* TAMARANG SA
 ROCURONIUM BROMIDE, ROCURONIUM BROMIDE

TARO

* TARO PHARMACEUTICAL INDUSTRIES LTD
 ACETAZOLAMIDE, ACETAZOLAMIDE
 AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE
 AZITHROMYCIN, AZITHROMYCIN
 BETAMETHASONE VALERATE, BETAMETHASONE VALERATE
 BROMPHENIRAMINE MALEATE, PSEUDOEPHEDRINE HYDROCHLORIDE AND DEXTROMETHORPHAN
 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)

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** T ****

- * TARO PHARMACEUTICAL INDUSTRIES LTD
 CHILDREN'S CETIRIZINE HYDROCHLORIDE HIVES RELIEF, CETIRIZINE HYDROCHLORIDE (OTC)
 CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE
 CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
 CLOMIPRAMINE HYDROCHLORIDE, CLOMIPRAMINE HYDROCHLORIDE
 CLORAZEPATE DIPOTASSIUM, CLORAZEPATE DIPOTASSIUM
 DESLORATADINE, DESLORATADINE
 DESONIDE, DESONIDE
 ENALAPRIL MALEATE, ENALAPRIL MALEATE
 ETODOLAC, ETODOLAC
 EXTENDED PHENYTOIN SODIUM, PHENYTOIN SODIUM
 FELBAMATE, FELBAMATE
 FLUCONAZOLE, FLUCONAZOLE
 FLUOROURACIL, FLUOROURACIL
 FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 GABAPENTIN, GABAPENTIN
 GRANISETRON HYDROCHLORIDE, GRANISETRON HYDROCHLORIDE
 HYDROCORTISONE BUTYRATE, HYDROCORTISONE BUTYRATE
 IMIQUIMOD, IMIQUIMOD
 KETOCONAZOLE, KETOCONAZOLE
 LAMOTRIGINE, LAMOTRIGINE
 LEVETIRACETAM, LEVETIRACETAM
 LORATADINE, LORATADINE (OTC)
 MELOXICAM, MELOXICAM
 METRONIDAZOLE, METRONIDAZOLE
 MINOXIDIL (FOR MEN), MINOXIDIL (OTC)
 MINOXIDIL (FOR WOMEN), MINOXIDIL (OTC)
 NORTRIPTYLINE HYDROCHLORIDE, NORTRIPTYLINE HYDROCHLORIDE
 NYSTATIN, NYSTATIN
 ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
 OXCARBAZEPINE, OXCARBAZEPINE
 PHENYTOIN, PHENYTOIN
 TERIL, CARBAMAZEPINE
 WARFARIN SODIUM, WARFARIN SODIUM
- * TARO PHARMACEUTICALS USA INC
 ACETIC ACID, ACETIC ACID, GLACIAL
 ACYCLOVIR, ACYCLOVIR
 ADAPALENE AND BENZOYL PEROXIDE, ADAPALENE
 ADAPALENE, ADAPALENE
 ALCLOMETASONE DIPROPIONATE, ALCLOMETASONE DIPROPIONATE
 AMMONIUM LACTATE, AMMONIUM LACTATE
 AZELAIC ACID, AZELAIC ACID
 BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
 CICLOPIROX, CICLOPIROX
 CLINDAMYCIN PHOSPHATE AND BENZOYL PEROXIDE, BENZOYL PEROXIDE
 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
 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)

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** T ****

* TARO PHARMACEUTICALS USA INC
 KETOZOLE, KETOCONAZOLE
 LIDOCAINE, LIDOCAINE
 LORATADINE, LORATADINE (OTC)
 MICONAZOLE 3, MICONAZOLE NITRATE (OTC)
 MOMETASONE FUROATE, MOMETASONE FUROATE
 MUPIROCIN, MUPIROCIN
 NYSTATIN AND TRIAMCINOLONE ACETONIDE, NYSTATIN
 NYSTATIN, NYSTATIN
 PHENYTOIN, PHENYTOIN
 PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
 RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
 RISPERIDONE, RISPERIDONE
 SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, SODIUM CHLORIDE
 STERILE WATER FOR INJECTION IN PLASTIC CONTAINER, STERILE WATER FOR INJECTION
 SULFACETAMIDE SODIUM, SULFACETAMIDE SODIUM
 TERBINAFINE HYDROCHLORIDE, TERBINAFINE HYDROCHLORIDE (OTC)
 TERCONAZOLE, TERCONAZOLE
 TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE
 TRIVAGIZOLE 3, CLOTRIMAZOLE (OTC)
 U-CORT, HYDROCORTISONE ACETATE

TARO PHARM INDS

* TARO PHARMACEUTICAL INDUSTRIES LTD
 AMCINONIDE, AMCINONIDE
 CARBAMAZEPINE, CARBAMAZEPINE
 CICLOPIROX, CICLOPIROX
 CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
 ENALAPRIL MALEATE AND HYDROCHLOROTHIAZIDE, ENALAPRIL MALEATE
 ETODOLAC, ETODOLAC
 HYDROCORTISONE BUTYRATE, HYDROCORTISONE BUTYRATE
 LAMOTRIGINE, LAMOTRIGINE

TARO PHARM INDS LTD

* TARO PHARMACEUTICAL INDUSTRIES LTD
 ACETAMINOPHEN, ACETAMINOPHEN (OTC)
 CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
 DESLORATADINE, DESLORATADINE
 FLUOCINONIDE EMULSIFIED BASE, FLUOCINONIDE
 HYDROCORTISONE AND ACETIC ACID, ACETIC ACID, GLACIAL
 HYDROCORTISONE, HYDROCORTISONE
 INFANTS' FEVERALL, ACETAMINOPHEN (OTC)
 LEVOCETIRIZINE DIHYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE
 TOPICORT, DESOXIMETASONE
 TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE

TARO PHARMS

* TARO PHARMACEUTICALS INC
 BUTENAFINE HYDROCHLORIDE, BUTENAFINE HYDROCHLORIDE (OTC)
 CLINDAMYCIN PHOSPHATE AND BENZOYL PEROXIDE, BENZOYL PEROXIDE
 DAPSONE, DAPSONE
 MICONAZOLE NITRATE, MICONAZOLE NITRATE (OTC)
 NAFTIFINE HYDROCHLORIDE, NAFTIFINE HYDROCHLORIDE
 OXICONAZOLE NITRATE, OXICONAZOLE NITRATE
 PLIAGLIS, LIDOCAINE
 TAZAROTENE, TAZAROTENE
 TOPICORT, DESOXIMETASONE
 TRETINOIN, TRETINOIN

TASMAN PHARMA

* TASMAN PHARMA INC
 CLOTRIMAZOLE, CLOTRIMAZOLE
 VERSACLOZ, CLOZAPINE

TCG FLUENT PHARMA

* TCG FLUENT PHARMA INVESTORS LP
 FLOLIPID, SIMVASTATIN

TEIKOKU PHARMA

* TEIKOKU PHARMA USA INC
 DOCETAXEL, DOCETAXEL

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** T ******TEIKOKU PHARMA USA**

* TEIKOKU PHARMA USA INC
LIDODERM, LIDOCAINE

TELIGENT

* TELIGENT OU
CEFOTAN, CEFOTETAN DISODIUM
CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE
FORTAZ, CEFTAZIDIME
ZANTAC, RANITIDINE HYDROCHLORIDE
ZINACEF, CEFUROXIME SODIUM

TELIGENT PHARMA INC

* TELIGENT PHARMA INC
BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
CICLOPIROX, CICLOPIROX
CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
CLOBETASOL PROPIONATE (EMOLLIENT), CLOBETASOL PROPIONATE
CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
DESONIDE, DESONIDE
DESOXIMETASONE, DESOXIMETASONE
DICLOFENAC SODIUM, DICLOFENAC SODIUM
DIFLORASONE DIACETATE, DIFLORASONE DIACETATE
ECONAZOLE NITRATE, ECONAZOLE NITRATE
ERYTHROMYCIN, ERYTHROMYCIN
FLUOCINONIDE, FLUOCINONIDE
FLURANDRENOLIDE, FLURANDRENOLIDE
GENTAMICIN SULFATE, GENTAMICIN SULFATE
HALOBETASOL PROPIONATE, HALOBETASOL PROPIONATE
HYDROCORTISONE BUTYRATE, HYDROCORTISONE BUTYRATE
HYDROCORTISONE, HYDROCORTISONE
LIDOCAINE AND PRILOCAINE, LIDOCAINE
LIDOCAINE HYDROCHLORIDE, LIDOCAINE HYDROCHLORIDE
LIDOCAINE, LIDOCAINE
NYSTATIN AND TRIAMCINOLONE ACETONIDE, NYSTATIN
TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE

TERSERA THERAPS LLC

* TERSERA THERAPEUTICS LLC
ERGOMAR, ERGOTAMINE TARTRATE
PRIALT, ZICONOTIDE ACETATE
QMIIZ ODT, MELOXICAM
VARUBI, ROLAPITANT HYDROCHLORIDE
ZOLADEX, GOSERELIN ACETATE

TETRAPHASE PHARMS

* TETRAPHASE PHARMACEUTICALS INC
XERAVA, ERAVACYCLINE DIHYDROCHLORIDE

TEVA

* TEVA NEUROSCIENCE INC
AZILECT, RASAGILINE MESYLATE

* TEVA PHARMACEUTICALS USA INC
ACETAMINOPHEN AND CODEINE PHOSPHATE, ACETAMINOPHEN
ACYCLOVIR, ACYCLOVIR
ADIPEX-P, PHENTERMINE HYDROCHLORIDE
ALBUTEROL SULFATE, ALBUTEROL SULFATE
AMOXICILLIN AND CLAVULANATE POTASSIUM, AMOXICILLIN
AMOXICILLIN PEDIATRIC, AMOXICILLIN
AMOXICILLIN, AMOXICILLIN
ATENOLOL, ATENOLOL
AZITHROMYCIN, AZITHROMYCIN
BENZAEPRILOL HYDROCHLORIDE, BENAZEPRIL HYDROCHLORIDE
BUSPIRONE HYDROCHLORIDE, BUSPIRONE HYDROCHLORIDE
CALCITRIOL, CALCITRIOL
CAPTOPRIL, CAPTOPRIL
CARVEDILOL, CARVEDILOL
CEFACLOR, CEFACLOL
CEFPROZIL, CEFPROZIL
CELECOXIB, CELECOXIB

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** T **

* TEVA PHARMACEUTICALS USA INC
 CEPHALEXIN, CEPHALEXIN
 CHILDREN'S FEXOFENADINE HYDROCHLORIDE ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
 CHILDREN'S FEXOFENADINE HYDROCHLORIDE HIVES, FEXOFENADINE HYDROCHLORIDE (OTC)
 CILOSTAZOL, CILOSTAZOL
 CIMETIDINE, CIMETIDINE
 CLARITHROMYCIN, CLARITHROMYCIN
 CLEMASTINE FUMARATE, CLEMASTINE FUMARATE
 CLONAZEPAM, CLONAZEPAM
 CLOPIDOGREL BISULFATE, CLOPIDOGREL BISULFATE
 CLOTRIMAZOLE, CLOTRIMAZOLE
 DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE
 DICLOFENAC POTASSIUM, DICLOFENAC POTASSIUM
 DICLOXACILLIN SODIUM, DICLOXACILLIN SODIUM
 DIFLUNISAL, DIFLUNISAL
 DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
 DISOPYRAMIDE PHOSPHATE, DISOPYRAMIDE PHOSPHATE
 DOXAZOSIN MESYLATE, DOXAZOSIN MESYLATE
 ENOXAPARIN SODIUM (PRESERVATIVE FREE), ENOXAPARIN SODIUM
 EPITOL, CARBAMAZEPINE
 ESZOPICLONE, ESZOPICLONE
 ETODOLAC, ETODOLAC
 FAMOTIDINE, FAMOTIDINE
 FAMOTIDINE, FAMOTIDINE (OTC)
 FEXOFENADINE HYDROCHLORIDE ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
 FEXOFENADINE HYDROCHLORIDE HIVES, FEXOFENADINE HYDROCHLORIDE (OTC)
 FEXOFENADINE HYDROCHLORIDE, FEXOFENADINE HYDROCHLORIDE
 FINASTERIDE, FINASTERIDE
 FLUCONAZOLE, FLUCONAZOLE
 FLUOCINONIDE EMULSIFIED BASE, FLUOCINONIDE
 FLUOCINONIDE, FLUOCINONIDE
 FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 FLURBIPROFEN, FLURBIPROFEN
 FLUVOXAMINE MALEATE, FLUVOXAMINE MALEATE
 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
 LORATADINE, LORATADINE (OTC)
 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 SODIUM, NAPROXEN SODIUM
 NAPROXEN, NAPROXEN
 NEFAZODONE HYDROCHLORIDE, NEFAZODONE HYDROCHLORIDE
 NEOMYCIN SULFATE, NEOMYCIN SULFATE
 NORTRIPTYLINE HYDROCHLORIDE, NORTRIPTYLINE HYDROCHLORIDE
 NYSTATIN, NYSTATIN
 OFLOXACIN, OFLOXACIN
 ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
 ONDANSETRON, ONDANSETRON
 OXAPROZIN, OXAPROZIN
 OXYMORPHONE HYDROCHLORIDE, OXYMORPHONE HYDROCHLORIDE
 PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** T ****

* TEVA PHARMACEUTICALS USA INC
 PAROXETINE HYDROCHLORIDE, PAROXETINE HYDROCHLORIDE
 PENICILLIN-VK, PENICILLIN V POTASSIUM
 PIROXICAM, PIROXICAM
 PRAVASTATIN SODIUM, PRAVASTATIN SODIUM
 PRELONE, PREDNISOLONE
 RIBAVIRIN, RIBAVIRIN
 SILDENAFIL CITRATE, SILDENAFIL CITRATE
 SOTALOL HYDROCHLORIDE, SOTALOL HYDROCHLORIDE
 SUCRALFATE, SUCRALFATE
 TICLOPIDINE HYDROCHLORIDE, TICLOPIDINE HYDROCHLORIDE
 TOLMETIN SODIUM, TOLMETIN SODIUM
 TOPIRAMATE, TOPIRAMATE
 TORSEMIDE, TORSEMIDE
 TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE
 VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE
 ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE

TEVA BRANDED PHARM

* TEVA BRANDED PHARMACEUTICAL PRODUCTS R AND D INC
 AUSTEDO, DEUTETRABENAZINE
 CONDYLOX, PODOFILOX
 EMLA, LIDOCAINE
 LOESTRIN 21 1.5/30, ETHINYL ESTRADIOL
 LOESTRIN 21 1/20, ETHINYL ESTRADIOL
 LOESTRIN 24 FE, ETHINYL ESTRADIOL
 LOESTRIN FE 1.5/30, ETHINYL ESTRADIOL
 LOESTRIN FE 1/20, ETHINYL ESTRADIOL
 LOSEASONIQUE, ETHINYL ESTRADIOL
 NOR-QD, NORETHINDRONE
 PROAIR DIGIHALER, ALBUTEROL SULFATE
 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

TEVA PARENTERAL

* TEVA PARENTERAL MEDICINES INC
 LEVALBUTEROL HYDROCHLORIDE, LEVALBUTEROL HYDROCHLORIDE

TEVA PHARM

* TEVA PHARMACEUTICAL INDUSTRIES LTD
 AIRDUO DIGIHALER, FLUTICASONE PROPIONATE
 AIRDUO RESPICLICK, FLUTICASONE PROPIONATE

TEVA PHARMS

* TEVA PHARMACEUTICALS USA
 AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE
 ANASTROZOLE, ANASTROZOLE
 AZITHROMYCIN, AZITHROMYCIN
 BISOPROLOL FUMARATE, BISOPROLOL FUMARATE
 BUDESONIDE, BUDESONIDE
 CARBAMAZEPINE, CARBAMAZEPINE
 CEFADROXIL, CEFADROXIL/CEFADROXIL HEMIHYDRATE
 CEFDINIR, CEFDINIR
 CEFPROZIL, CEFPROZIL
 CETIRIZINE HYDROCHLORIDE, CETIRIZINE HYDROCHLORIDE
 CROMOLYN SODIUM, CROMOLYN SODIUM
 DORZOLAMIDE HYDROCHLORIDE AND TIMOLOL MALEATE, DORZOLAMIDE HYDROCHLORIDE
 DOXEPIN HYDROCHLORIDE, DOXEPIN HYDROCHLORIDE
 FAMCICLOVIR, FAMCICLOVIR
 FLUVASTATIN SODIUM, FLUVASTATIN SODIUM
 GABAPENTIN, GABAPENTIN
 GEMCITABINE HYDROCHLORIDE, GEMCITABINE HYDROCHLORIDE
 GLIPIZIDE AND METFORMIN HYDROCHLORIDE, GLIPIZIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** T ****

* TEVA PHARMACEUTICALS USA
 HYDROCORTISONE, HYDROCORTISONE
 HYDROXYCHLOROQUINE SULFATE, HYDROXYCHLOROQUINE SULFATE
 IRBESARTAN, IRBESARTAN
 LANSOPRAZOLE, LANSOPRAZOLE
 LEFLUNOMIDE, LEFLUNOMIDE
 LETROZOLE, LETROZOLE
 LEVETIRACETAM, LEVETIRACETAM
 LEVOCETIRIZINE DIHYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE
 LINEZOLID, LINEZOLID
 LOSARTAN POTASSIUM AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 MEGESTROL ACETATE, MEGESTROL ACETATE
 MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
 METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
 MONTELUKAST SODIUM, MONTELUKAST SODIUM
 MYCOPHENOLATE MOFETIL, MYCOPHENOLATE MOFETIL
 OLANZAPINE AND FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 OXALIPLATIN, OXALIPLATIN
 PACLITAXEL, PACLITAXEL
 PRAVASTATIN SODIUM, PRAVASTATIN SODIUM
 PRAZOSIN HYDROCHLORIDE, PRAZOSIN HYDROCHLORIDE
 PREGABALIN, PREGABALIN
 QUETIAPINE FUMARATE, QUETIAPINE FUMARATE
 QUININE SULFATE, QUININE SULFATE
 RAMIPRIL, RAMIPRIL
 RIZATRIPTAN BENZOATE, RIZATRIPTAN BENZOATE
 ROCURONIUM BROMIDE, ROCURONIUM BROMIDE
 SILDENAFIL CITRATE, SILDENAFIL CITRATE
 TAMSULOSIN HYDROCHLORIDE, TAMSULOSIN HYDROCHLORIDE
 TRANDOLAPRIL, TRANDOLAPRIL
 URSODIOL, URSODIOL
 VANDAZOLE, METRONIDAZOLE
 VARDENAFIL HYDROCHLORIDE, VARDENAFIL HYDROCHLORIDE
 ZALEPLON, ZALEPLON

TEVA PHARMS INTL

* TEVA PHARMACEUTICALS INTERNATIONAL GMBH
 AMRIX, CYCLOBENZAPRINE HYDROCHLORIDE
 SYNRIPO, OMACETAXINE MEPESUCCINATE

TEVA PHARMS USA

* TEVA PHARMACEUTICALS USA
 ABACAVIR SULFATE AND LAMIVUDINE, ABACAVIR SULFATE
 ACITRETIN, ACITRETIN
 ADENOSINE, ADENOSINE
 ALMOTRIPTAN MALATE, ALMOTRIPTAN MALATE
 ALPROSTADIL, ALPROSTADIL
 ARGATROBAN IN 0.9% SODIUM CHLORIDE, ARGATROBAN
 ATAZANAVIR SULFATE, ATAZANAVIR SULFATE
 ATOMOXETINE HYDROCHLORIDE, ATOMOXETINE HYDROCHLORIDE
 ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM
 BLEOMYCIN SULFATE, BLEOMYCIN SULFATE
 BUDESONIDE, BUDESONIDE
 CARBOPLATIN, CARBOPLATIN
 CLARAVIS, ISOTRETINOIN
 CLOBAZAM, CLOBAZAM
 CLOZAPINE, CLOZAPINE
 COPAXONE, GLATIRAMER ACETATE
 DACARBAZINE, DACARBAZINE
 DAUNORUBICIN HYDROCHLORIDE, DAUNORUBICIN HYDROCHLORIDE
 DESMOPRESSIN ACETATE, DESMOPRESSIN ACETATE
 DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE
 DEXMETHYLPHENIDATE HYDROCHLORIDE, DEXMETHYLPHENIDATE HYDROCHLORIDE
 DOXORUBICIN HYDROCHLORIDE, DOXORUBICIN HYDROCHLORIDE
 ENTECAVIR, ENTECAVIR
 EPOPROSTENOL SODIUM, EPOPROSTENOL SODIUM
 EPTIFIBATIDE, EPTIFIBATIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** T ****

- * TEVA PHARMACEUTICALS USA
ERLOTINIB HYDROCHLORIDE, ERLOTINIB HYDROCHLORIDE
ESCITALOPRAM OXALATE, ESCITALOPRAM OXALATE
ESTRADIOL, ESTRADIOL
ETOPOSIDE, ETOPOSIDE
EZETIMIBE, EZETIMIBE
FLUOROURACIL, FLUOROURACIL
FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
FLUVASTATIN SODIUM, FLUVASTATIN SODIUM
FOSAPREPITANT DIMEGLUMINE, FOSAPREPITANT DIMEGLUMINE
GABAPENTIN, GABAPENTIN
GUANFACINE HYDROCHLORIDE, GUANFACINE HYDROCHLORIDE
HALOPERIDOL DECANOATE, HALOPERIDOL DECANOATE
HALOPERIDOL, HALOPERIDOL LACTATE
IDARUBICIN HYDROCHLORIDE PFS, IDARUBICIN HYDROCHLORIDE
IFOSFAMIDE, IFOSFAMIDE
IMATINIB MESYLATE, IMATINIB MESYLATE
IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE
LANSOPRAZOLE, LANSOPRAZOLE
LEUCOVORIN CALCIUM, LEUCOVORIN CALCIUM
LEUPROLIDE ACETATE, LEUPROLIDE ACETATE
LEVALBUTEROL HYDROCHLORIDE, LEVALBUTEROL HYDROCHLORIDE
LINEZOLID, LINEZOLID
LOGILIA, ULIPRISTAL ACETATE
MESNA, MESNA
METHYLPREDNISOLONE ACETATE, METHYLPREDNISOLONE ACETATE
METOCLOPRAMIDE HYDROCHLORIDE, METOCLOPRAMIDE HYDROCHLORIDE
MITOXANTRONE HYDROCHLORIDE, MITOXANTRONE HYDROCHLORIDE
MORPHINE SULFATE, MORPHINE SULFATE
MOXIFLOXACIN HYDROCHLORIDE, MOXIFLOXACIN HYDROCHLORIDE
MYCOPHENOLIC ACID, MYCOPHENOLIC ACID
NOREPINEPHRINE BITARTRATE, NOREPINEPHRINE BITARTRATE
OCTREOTIDE ACETATE, OCTREOTIDE ACETATE
OMEGA-3-ACID ETHYL ESTERS, OMEGA-3-ACID ETHYL ESTERS
OMEPRAZOLE, OMEPRAZOLE
PALONOSETRON HYDROCHLORIDE, PALONOSETRON HYDROCHLORIDE
PAMIDRONATE DISODIUM, PAMIDRONATE DISODIUM
PARICALCITOL, PARICALCITOL
PIOGLITAZONE HYDROCHLORIDE AND METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
PIOGLITAZONE HYDROCHLORIDE, PIOGLITAZONE HYDROCHLORIDE
RALOXIFENE HYDROCHLORIDE, RALOXIFENE HYDROCHLORIDE
RISEDRONATE SODIUM, RISEDRONATE SODIUM
SOLIFENACIN SUCCINATE, SOLIFENACIN SUCCINATE
TADALAFIL, TADALAFIL
TENOFVIR DISOPROXIL FUMARATE, TENOFVIR DISOPROXIL FUMARATE
TOBRAMYCIN SULFATE, TOBRAMYCIN SULFATE
TOBRAMYCIN, TOBRAMYCIN
TOLTERODINE TARTRATE, TOLTERODINE TARTRATE
TOPOTECAN HYDROCHLORIDE, TOPOTECAN HYDROCHLORIDE
TREPROSTINIL, TREPROSTINIL
VECURONIUM BROMIDE, VECURONIUM BROMIDE
VIGABATRIN, VIGABATRIN
VILAZODONE HYDROCHLORIDE, VILAZODONE HYDROCHLORIDE
VINCRISTINE SULFATE PFS, VINCRISTINE SULFATE
VINORELBINE TARTRATE, VINORELBINE TARTRATE
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
ELETRIPTAN HYDROBROMIDE, ELETRIPTAN HYDROBROMIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** T ****

* TEVA PHARMACEUTICALS USA INC
 EMTRICITABINE AND TENOFOVIR DISOPROXIL FUMARATE, EMTRICITABINE
 EPINEPHRINE (AUTOINJECTOR), EPINEPHRINE
 ESTRADIOL, ESTRADIOL
 EVEROLIMUS, EVEROLIMUS
 ICATIBANT ACETATE, ICATIBANT ACETATE
 IVERMECTIN, IVERMECTIN
 LIDOCAINE, LIDOCAINE
 MEDROXYPROGESTERONE ACETATE, MEDROXYPROGESTERONE ACETATE
 MESALAMINE, MESALAMINE
 METHYLERGONOVINE MALEATE, METHYLERGONOVINE MALEATE
 METRONIDAZOLE, METRONIDAZOLE
 OLMESARTAN MEDOXMIL AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 OSELTAMIVIR PHOSPHATE, OSELTAMIVIR PHOSPHATE
 OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE
 PENICILLAMINE, PENICILLAMINE
 RABEPRAZOLE SODIUM, RABEPRAZOLE SODIUM
 SELFEMRA, FLUOXETINE HYDROCHLORIDE
 TRAZODONE HYDROCHLORIDE, TRAZODONE HYDROCHLORIDE
 TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE

TEVA PHARMS USA INC

* TEVA PHARMACEUTICALS USA INC
 FULVESTRANT, FULVESTRANT
 POTASSIUM CITRATE, POTASSIUM CITRATE

THE FEINSTEIN INST

* THE FEINSTEIN INSTITUTE FOR MEDICAL RESEARCH
 SODIUM FLUORIDE F-18, SODIUM FLUORIDE F-18

THEPHARMANETWORK LLC

* THEPHARMANETWORK LLC
 ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM
 BENZONATATE, BENZONATATE
 ISONIAZID, ISONIAZID
 METHADONE HYDROCHLORIDE, METHADONE HYDROCHLORIDE
 NIMODIPINE, NIMODIPINE
 THERMAZENE, SILVER SULFADIAZINE

THERAPEUTICSMD INC

* THERAPEUTICSMD INC
 ANNOVERA, ETHINYL ESTRADIOL
 BIJUVA, ESTRADIOL
 IMVEXXY, ESTRADIOL

THERATECHNOLOGIES

* THERATECHNOLOGIES INC
 EGRIFTA, TESAMORELIN ACETATE

TIANJIN TIANYAO

* TIANJIN TIANYAO PHARMACEUTICALS CO LTD
 CELECOXIB, CELECOXIB
 METHYLPREDNISOLONE, METHYLPREDNISOLONE

TIME-CAP LABS INC

* TIME-CAP LABORATORIES INC
 VALACYCLOVIR HYDROCHLORIDE, VALACYCLOVIR HYDROCHLORIDE

TITAN PHARMS

* TITAN PHARMACEUTICALS INC
 PROBUPHINE, BUPRENORPHINE HYDROCHLORIDE

TOLMAR

* TOLMAR INC
 ACYCLOVIR, ACYCLOVIR
 ADAPALENE AND BENZOYL PEROXIDE, ADAPALENE
 ADAPALENE, ADAPALENE
 ATRIDOX, DOXYCYCLINE HYCLATE
 AZELAIC ACID, AZELAIC ACID
 CALCIPOTRIENE AND BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
 CALCIPOTRIENE, CALCIPOTRIENE
 CLINDAMYCIN PHOSPHATE AND BENZOYL PEROXIDE, BENZOYL PEROXIDE
 CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
 DICLOFENAC SODIUM, DICLOFENAC SODIUM

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** T ****

* TOLMAR INC
 ERYTHROMYCIN AND BENZOYL PEROXIDE, BENZOYL PEROXIDE
 KETOCONAZOLE, KETOCONAZOLE
 LIDOCAINE AND PRILOCAINE, LIDOCAINE
 METRONIDAZOLE, METRONIDAZOLE
 NAFTIFINE HYDROCHLORIDE, NAFTIFINE HYDROCHLORIDE

TOLMAR THERAP

* TOLMAR THERAPEUTICS INC
 ELIGARD, LEUPROLIDE ACETATE

TOPROL

* TOPROL ACQUISITION LLC
 TOPROL-XL, METOPROLOL SUCCINATE
 ZONTIVITY, VORAPAXAR SULFATE

TORPHARM

* TORPHARM INC
 CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE

TORRENT

* TORRENT PHARMA INC
 MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE

* TORRENT PHARMACEUTICALS LTD
 ACYCLOVIR, ACYCLOVIR
 AMLODIPINE AND OLMESARTAN MEDOXOMIL, AMLODIPINE BESYLATE
 ANAGRELIDE HYDROCHLORIDE, ANAGRELIDE HYDROCHLORIDE
 ARIPIPRAZOLE, ARIPIPRAZOLE
 BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
 CELECOXIB, CELECOXIB
 CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
 DARIFENACIN HYDROBROMIDE, DARIFENACIN HYDROBROMIDE
 DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE
 ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM
 FENOFIBRATE (MICRONIZED), FENOFIBRATE
 FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 FLUVOXAMINE MALEATE, FLUVOXAMINE MALEATE
 ITRACONAZOLE, ITRACONAZOLE
 LAMOTRIGINE, LAMOTRIGINE
 MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 MOMETASONE FUROATE, MOMETASONE FUROATE
 MONTELUKAST SODIUM, MONTELUKAST SODIUM
 MOXIFLOXACIN HYDROCHLORIDE, MOXIFLOXACIN HYDROCHLORIDE
 NYSTATIN, NYSTATIN
 OLMESARTAN MEDOXOMIL AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 OLMESARTAN MEDOXOMIL, AMLODIPINE AND HYDROCHLOROTHIAZIDE, AMLODIPINE BESYLATE
 OLMESARTAN MEDOXOMIL, OLMESARTAN MEDOXOMIL
 RABEPRAZOLE SODIUM, RABEPRAZOLE SODIUM
 ROFLUMILAST, ROFLUMILAST
 ROSUVASTATIN CALCIUM, ROSUVASTATIN CALCIUM
 SILDENAFIL CITRATE, SILDENAFIL CITRATE
 TADALAFIL, TADALAFIL
 TELMISARTAN AND AMLODIPINE, AMLODIPINE BESYLATE
 TELMISARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 TELMISARTAN, TELMISARTAN
 TOLTERODINE TARTRATE, TOLTERODINE TARTRATE
 TRAZODONE HYDROCHLORIDE, TRAZODONE HYDROCHLORIDE

TORRENT PHARMS

* TORRENT PHARMACEUTICALS LIMITED
 LEVOFLOXACIN, LEVOFLOXACIN

* TORRENT PHARMACEUTICALS LTD
 AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
 CARBAMAZEPINE, CARBAMAZEPINE
 CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE
 DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
 ISOSORBIDE MONONITRATE, ISOSORBIDE MONONITRATE
 LAMOTRIGINE, LAMOTRIGINE
 LEVETIRACETAM, LEVETIRACETAM

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** T ****

- * TORRENT PHARMACEUTICALS LTD
 LOSARTAN POTASSIUM AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 LOSARTAN POTASSIUM, LOSARTAN POTASSIUM
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
 PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE
 RISPERIDONE, RISPERIDONE
 SERTRALINE HYDROCHLORIDE, SERTRALINE HYDROCHLORIDE
 TOPIRAMATE, TOPIRAMATE
 ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE
- * TORRENT PHARMACEUTICALS LTD.
 ALFUZOSIN HYDROCHLORIDE, ALFUZOSIN HYDROCHLORIDE
- TORRENT PHARMS LLC**
- * TORRENT PHARMACEUTICALS LLC
 CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 CETIRIZINE HYDROCHLORIDE HIVES RELIEF, CETIRIZINE HYDROCHLORIDE (OTC)
 OLANZAPINE, OLANZAPINE
 VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE
- TORRENT PHARMS LTD**
- * TORRENT PHARMACEUTICALS LTD
 CLOPIDOGREL BISULFATE, CLOPIDOGREL BISULFATE
 ESCITALOPRAM OXALATE, ESCITALOPRAM OXALATE
 FELODIPINE, FELODIPINE
 LEVETIRACETAM, LEVETIRACETAM
 MONTELUKAST SODIUM, MONTELUKAST SODIUM
 OLANZAPINE, OLANZAPINE
 PIOGLITAZONE HYDROCHLORIDE AND METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 PIOGLITAZONE HYDROCHLORIDE, PIOGLITAZONE HYDROCHLORIDE
 QUETIAPINE FUMARATE, QUETIAPINE FUMARATE
 SILDENAFIL CITRATE, SILDENAFIL CITRATE
- TRIS PHARMA INC**
- * TRIS PHARMA INC
 CHILDREN'S CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 CHILDREN'S CETIRIZINE HYDROCHLORIDE HIVES RELIEF, CETIRIZINE HYDROCHLORIDE (OTC)
 DEXMETHYLPHENIDATE HYDROCHLORIDE, DEXMETHYLPHENIDATE HYDROCHLORIDE
 DEXTROAMPHETAMINE SULFATE, DEXTROAMPHETAMINE SULFATE
 DEXTROMETHORPHAN POLISTIREX, DEXTROMETHORPHAN POLISTIREX (OTC)
 DYANAVEL XR, AMPHETAMINE
 GABAPENTIN, GABAPENTIN
 HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
 HYDROCODONE POLISTIREX AND CHLORPHENIRAMINE POLISTIREX, CHLORPHENIRAMINE POLISTIREX
 IBUPROFEN, IBUPROFEN (OTC)
 LEVETIRACETAM, LEVETIRACETAM
 METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
 MORPHINE SULFATE, MORPHINE SULFATE
 POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
 PROMETHAZINE HYDROCHLORIDE AND CODEINE PHOSPHATE, CODEINE PHOSPHATE
 PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
 RISPERIDONE, RISPERIDONE
 THEOPHYLLINE, THEOPHYLLINE
- TRUSTEES UNIV PA**
- * TRUSTEES OF THE UNIV OF PENNSYLVANIA
 FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18
- TULEX PHARMS INC**
- * TULEX PHARMACEUTICALS INC
 OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE
- TWI PHARMS**
- * TWI PHARMACEUTICALS INC
 BENZPHETAMINE HYDROCHLORIDE, BENZPHETAMINE HYDROCHLORIDE
 BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
 CYPROHEPTADINE HYDROCHLORIDE, CYPROHEPTADINE HYDROCHLORIDE
 DICLOFENAC SODIUM, DICLOFENAC SODIUM
 DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
 DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
 GUANFACINE HYDROCHLORIDE, GUANFACINE HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** T ****

* TWI PHARMACEUTICALS INC
 HYDROXYCHLOROQUINE SULFATE, HYDROXYCHLOROQUINE SULFATE
 LABETALOL HYDROCHLORIDE, LABETALOL HYDROCHLORIDE
 MEGESTROL ACETATE, MEGESTROL ACETATE
 METOPROLOL SUCCINATE, METOPROLOL SUCCINATE
 NIFEDIPINE, NIFEDIPINE
 SEVELAMER CARBONATE, SEVELAMER CARBONATE
 TESTOSTERONE, TESTOSTERONE
 ZOLMITRIPTAN, ZOLMITRIPTAN

TWI PHARMS INC

* TWI PHARMACEUTICALS INC
 CYCLOBENZAPRINE HYDROCHLORIDE, CYCLOBENZAPRINE HYDROCHLORIDE

**** U ******UBI**

* UBI PHARMA INC
 DESMOPRESSIN ACETATE, DESMOPRESSIN ACETATE

UCB INC

* UCB INC
 BRIVIACT, BRIVARACETAM
 KEPPRA XR, LEVETIRACETAM
 KEPPRA, LEVETIRACETAM
 NAYZILAM, MIDAZOLAM
 NEUPRO, ROTIGOTINE
 VIMPAT, LACOSAMIDE

UCLA BIOMEDICAL

* UCLA BIOMEDICAL CYCLOTRON
 AMMONIA N 13, AMMONIA N-13
 FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18

UCSF RODIOPHARM

* UCSF RADIOPHARMACEUTICAL 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

UIHC PET IMAGING

* UNIV IOWA HOSPS AND CLINICS PET IMAGING CENTER
 FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18
 GALLIUM DOTATOC GA 68, GALLIUM DOTATOC GA-68

ULTRATAB LABS INC

* ULTRATAB LABORATORIES INC
 IBUPROFEN, IBUPROFEN (OTC)

UMEDICA LABS PVT LTD

* UMEDICA LABORATORIES PRIVATE LTD
 CELECOXIB, CELECOXIB
 CHLORTHALIDONE, CHLORTHALIDONE
 OLMESARTAN MEDOXOMIL AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 OLMESARTAN MEDOXOMIL, OLMESARTAN MEDOXOMIL

UNICHEM

* UNICHEM LABORATORIES LTD
 BISOPROLOL FUMARATE AND HYDROCHLOROTHIAZIDE, BISOPROLOL FUMARATE
 CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 CETIRIZINE HYDROCHLORIDE HIVES, CETIRIZINE HYDROCHLORIDE (OTC)
 CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
 HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 MELOXICAM, MELOXICAM
 ZALEPLON, ZALEPLON

UNICHEM LABS LTD

* UNICHEM LABORATORIES LIMITED
 DIVALPROEX SODIUM, DIVALPROEX SODIUM

* UNICHEM LABORATORIES LTD
 ALFUZOSIN HYDROCHLORIDE, ALFUZOSIN HYDROCHLORIDE
 ALLOPURINOL, ALLOPURINOL
 AMLODIPINE BESYLATE, AMLODIPINE BESYLATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** U ******* UNICHEM LABORATORIES LTD**

ATENOLOL, ATENOLOL
 BUSPIRONE HYDROCHLORIDE, BUSPIRONE HYDROCHLORIDE
 CHLORTHALIDONE, CHLORTHALIDONE
 DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
 IRBESARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 IRBESARTAN, IRBESARTAN
 LAMOTRIGINE, LAMOTRIGINE
 LOSARTAN POTASSIUM AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 LOSARTAN POTASSIUM, LOSARTAN POTASSIUM
 MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
 METRONIDAZOLE, METRONIDAZOLE
 MONTELUKAST SODIUM, MONTELUKAST SODIUM
 PIROXICAM, PIROXICAM
 PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE
 QUETIAPINE FUMARATE, QUETIAPINE FUMARATE
 RIZATRIPTAN BENZOATE, RIZATRIPTAN BENZOATE
 SOLIFENACIN SUCCINATE, SOLIFENACIN SUCCINATE
 TADALAFIL, TADALAFIL
 TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE
 TOPIRAMATE, TOPIRAMATE
 TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE
 VALSARTAN, VALSARTAN

UNICHEM PHARMS (USA)

* UNICHEM PHARMACEUTICALS (USA) INC
 BISOPROLOL FUMARATE, BISOPROLOL FUMARATE

UNIMARK REMEDIES LTD

* UNIMARK REMEDIES LTD
 MONTELUKAST SODIUM, MONTELUKAST SODIUM

UNIQUE PHARM LABS

* UNIQUE PHARMACEUTICAL LABORATORIES A DIVISION OF J.B. CHEMICALS AND PHARMACEUTICALS LTD
 ATENOLOL, ATENOLOL
 CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 CETIRIZINE HYDROCHLORIDE HIVES, CETIRIZINE HYDROCHLORIDE (OTC)
 CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE
 DICLOFENAC SODIUM, DICLOFENAC SODIUM
 FEXOFENADINE HYDROCHLORIDE ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
 FLUCONAZOLE, FLUCONAZOLE
 GLIPIZIDE, GLIPIZIDE
 LITHIUM CARBONATE, LITHIUM CARBONATE
 MIDODRINE HYDROCHLORIDE, MIDODRINE HYDROCHLORIDE
 OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE
 RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE (OTC)
 TINIDAZOLE, TINIDAZOLE
 TOLTERODINE TARTRATE, TOLTERODINE TARTRATE

UNITED BIOMEDCL

* UNITED BIOMEDICAL INC
 TERBUTALINE SULFATE, TERBUTALINE SULFATE

UNITED GUARDIAN

* UNITED GUARDIAN INC
 RENACIDIN, CITRIC ACID

UNITED THERAP

* UNITED THERAPEUTICS CORP
 ORENITRAM, TREPROSTINIL DIOLAMINE
 REMODULIN, TREPROSTINIL
 TYVASO, TREPROSTINIL

UNIV MICHIGAN

* UNIV MICHIGAN PET RADIOPHARMACEUTICAL PRODUCTION PROGRAM
 FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18

UNIV TX MD ANDERSON

* UNIV TEXAS MD ANDERSON CANCER CENTER
 CHOLINE C-11, CHOLINE C-11
 FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18

UNIV TX SW MEDCTR

* UNIV TEXAS SOUTHWESTERN MEDCTR

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** U ****

* UNIV TEXAS SOUTHWESTERN MEDCTR
 AMMONIA N 13, AMMONIA N-13
 FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18

UNIV UTAH CYCLOTRON

* UNIV UTAH CYCLOTRON RADIOCHEMISTRY LAB
 FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18
 SODIUM FLUORIDE F-18, SODIUM FLUORIDE F-18

UPSHER SMITH LABS

* UPSHER SMITH LABORATORIES INC
 FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
 MORPHINE SULFATE, MORPHINE SULFATE

* UPSHER SMITH LABORATORIES LLC
 AMANTADINE HYDROCHLORIDE, AMANTADINE HYDROCHLORIDE
 AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
 AZELASTINE HYDROCHLORIDE, AZELASTINE HYDROCHLORIDE
 BACLOFEN, BACLOFEN
 BETHANECHOL CHLORIDE, BETHANECHOL CHLORIDE
 BEXAROTENE, BEXAROTENE
 BUMETANIDE, BUMETANIDE
 CHLORPROMAZINE HYDROCHLORIDE, CHLORPROMAZINE HYDROCHLORIDE
 CLOBAZAM, CLOBAZAM
 CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
 DIPHENOXYLATE HYDROCHLORIDE AND ATROPINE SULFATE, ATROPINE SULFATE
 DIVALPROEX SODIUM, DIVALPROEX SODIUM
 DOXAZOSIN MESYLATE, DOXAZOSIN MESYLATE
 ETHACRYNIC ACID, ETHACRYNIC ACID
 EXEMESTANE, EXEMESTANE
 FLUOXYMESTERONE, FLUOXYMESTERONE
 FLUVOXAMINE MALEATE, FLUVOXAMINE MALEATE
 FOSINOPRIL SODIUM, FOSINOPRIL SODIUM
 JANTOVEN, WARFARIN SODIUM
 KLOL-CON M10, POTASSIUM CHLORIDE
 KLOL-CON M15, POTASSIUM CHLORIDE
 KLOL-CON M20, POTASSIUM CHLORIDE
 KLOL-CON, POTASSIUM CHLORIDE
 LOSARTAN POTASSIUM, LOSARTAN POTASSIUM
 MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
 MIRTAZAPINE, MIRTAZAPINE
 MORPHINE SULFATE, MORPHINE SULFATE
 NYSTATIN, NYSTATIN
 ORVATEN, MIDODRINE HYDROCHLORIDE
 OXANDROLONE, OXANDROLONE
 OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE
 PACERONE, AMIODARONE HYDROCHLORIDE
 PENTOXIL, PENTOXIFYLLINE
 PREVALITE, CHOLESTYRAMINE
 QUDEXY XR, TOPIRAMATE
 SORINE, SOTALOL HYDROCHLORIDE
 TOPIRAMATE, TOPIRAMATE
 TOSYMRA, SUMATRIPTAN
 VOGELXO, TESTOSTERONE
 ZEMBRACE SYMTOUCH, SUMATRIPTAN SUCCINATE

US PHARM HOLDINGS

* US PHARMACEUTICAL HOLDINGS II LLC
 DRISDOL, ERGOCALCIFEROL
 HIPREX, METHENAMINE HIPPURATE
 LASIX, FUROSEMIDE
 NORPRAMIN, DESIPRAMINE HYDROCHLORIDE

US PHARMS HOLDINGS I

* US PHARMACEUTICALS HOLDINGS I LLC
 LOPRESSOR HCT, HYDROCHLOROTHIAZIDE
 LOPRESSOR, METOPROLOL TARTRATE
 LOTENSIN HCT, BENAZEPRIL HYDROCHLORIDE
 LOTENSIN, BENAZEPRIL HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** U ****

* US PHARMACEUTICALS HOLDINGS I LLC
PARLODEL, BROMOCRIPTINE MESYLATE

US WORLDMEDS

* US WORLDMEDS LLC
APOKYN, APOMORPHINE HYDROCHLORIDE
REVONTO, DANTROLENE SODIUM

US WORLDMEDS LLC

* US WORLDMEDS LLC
CORGARD, NADOLOL
LUCEMYRA, LOFEXIDINE HYDROCHLORIDE
XADAGO, SAFINAMIDE MESYLATE

USPHARMA

* USPHARMA LTD
NITRO-DUR, NITROGLYCERIN

USPHARMA WINDLAS

* USPHARMA WINDLAS LLC
PRASUGREL, PRASUGREL HYDROCHLORIDE

USV

* USV PRIVATE LTD
OLOPATADINE HYDROCHLORIDE, OLOPATADINE HYDROCHLORIDE
ZOLEDRONIC ACID, ZOLEDRONIC ACID

**** V ******VALEANT**

* VALEANT PHARMACEUTICALS INTERNATIONAL
BONTRIL PDM, PHENDIMETRAZINE TARTRATE
MYSOLINE, PRIMIDONE

VALEANT BERMUDA

* VALEANT INTERNATIONAL BERMUDA
DERMATOP E EMOLLIENT, PREDNICARBATE
PENLAC, CICLOPIROX
RETIN-A, TRETINOIN

VALEANT INTL

* VALEANT INTERNATIONAL BARBADOS SRL
RETIN-A MICRO, TRETINOIN
RETIN-A, TRETINOIN
RETIN-A-MICRO, TRETINOIN
WELLBUTRIN XL, BUPROPION HYDROCHLORIDE
* VALEANT INTERNATIONAL SRL
BENZAMYCIN, BENZOYL PEROXIDE

VALEANT LUXEMBOURG

* VALEANT PHARMACEUTICALS LUXEMBOURG SARL
TARGRETIN, BEXAROTENE
VISUDYNE, VERTEPORFIN

VALEANT PHARM INTL

* VALEANT PHARMACEUTICALS INTERNATIONAL
ANDROID 25, METHYLTESTOSTERONE
LIBRIUM, CHLORDIAZEPOXIDE HYDROCHLORIDE
TESTRED, METHYLTESTOSTERONE
VIRAZOLE, RIBAVIRIN

VALEANT PHARMS

* VALEANT PHARMACEUTICALS NORTH AMERICA LLC
LIBRAX, CHLORDIAZEPOXIDE HYDROCHLORIDE
PENTOXIFYLLINE, PENTOXIFYLLINE

VALEANT PHARMS INC

* VALEANT PHARMACEUTICALS INTERNATIONAL INC
GRIS-PEG, GRISEOFULVIN, ULTRAMICROSIZE

VALEANT PHARMS INTL

* VALEANT PHARMACEUTICALS INTERNATIONAL
APRISO, MESALAMINE
COLAZAL, BALSALAZIDE DISODIUM
CUPRIMINE, PENICILLAMINE
LACRISERT, HYDROXYPROPYL CELLULOSE
TIMOPTIC IN OCUDOSE, TIMOLOL MALEATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** V ****

* VALEANT PHARMACEUTICALS INTERNATIONAL
TIMOPTIC, TIMOLOL MALEATE

VALEANT PHARMS LLC

* VALEANT PHARMACEUTICALS NORTH AMERICA LLC
MACUGEN, PEGAPTANIB SODIUM
TIMOPTIC-XE, TIMOLOL MALEATE

VALEANT PHARMS NORTH

* VALEANT PHARMACEUTICALS NORTH AMERICA LLC
APLENZIN, BUPROPION HYDROBROMIDE
CARAC, FLUOROURACIL
DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
FENOFIBRATE, FENOFIBRATE
KLARON, SULFACETAMIDE SODIUM
NIFEDIPINE, NIFEDIPINE
NORITATE, METRONIDAZOLE
RENOVA, TRETINOIN
RETIN-A, TRETINOIN
SECONAL SODIUM, SECOBARBITAL SODIUM
VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE
XENAZINE, TETRABENAZINE

VALIDUS PHARMS

* VALIDUS PHARMACEUTICALS LLC
BUMEX, BUMETANIDE
EQUETRO, CARBAMAZEPINE
ROCALTROL, CALCITRIOL

VALIDUS PHARMS INC

* VALIDUS PHARMACEUTICALS INC
MARPLAN, ISOCARBOXAZID

VANDA PHARMS INC

* VANDA PHARMACEUTICALS INC
FANAPT, ILOPERIDONE
HETLIOZ, TASIMELTEON

VELOXIS PHARMS INC

* VELOXIS PHARMACEUTICALS INC
ENVARUSUS XR, TACROLIMUS

VERASTEM INC

* VERASTEM INC
COPIKTRA, DUVELISIB

VERO BIOTECH

* VERO BIOTECH
GENOSYL, NITRIC OXIDE

VEROSCIENCE

* VEROSCIENCE LLC
CYCLOSET, BROMOCRIPTINE MESYLATE

VERTEX PHARMS

* VERTEX PHARMACEUTICALS INC
KALYDECO, IVACAFTOR

VERTEX PHARMS INC

* VERTEX PHARMACEUTICALS INC
KALYDECO, IVACAFTOR
ORKAMBI, IVACAFTOR
SYMDEKO (COPACKAGED), IVACAFTOR
TRIKAFTA (COPACKAGED), ELEXACAFTOR, IVACAFTOR, TEZACAFTOR

VERTICAL PHARMS LLC

* VERTICAL PHARMACEUTICALS LLC
DIVIGEL, ESTRADIOL

VGYAAN

* VGYAAN PHARMACEUTICALS LLC
NADOLOL, NADOLOL

VICURON

* VICURON PHARMACEUTICALS INC
ERAXIS, ANIDULAFUNGIN

VIFOR FRESENIUS

* VIFOR FRESENIUS MEDICAL CARE RENAL PHARMA FRANCE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** V ****

* VIFOR FRESENIUS MEDICAL CARE RENAL PHARMA FRANCE
VELPHORO, SUCROFERRIC OXYHYDROXIDE

VIIIV HLTHCARE

* VIIIV HEALTHCARE CO
COMBIVIR, LAMIVUDINE
DOVATO, DOLUTEGRAVIR SODIUM
EPIVIR, LAMIVUDINE
EPZICOM, ABACAVIR SULFATE
JULUCA, DOLUTEGRAVIR SODIUM
LEXIVA, FOSAMPRENAVIR CALCIUM
RESCRIPTOR, DELAVIRDINE MESYLATE
RETROVIR, ZIDOVUDINE
SELZENTRY, MARAVIROC
TIVICAY, DOLUTEGRAVIR SODIUM
TRIUMEQ, ABACAVIR SULFATE
TRIZIVIR, ABACAVIR SULFATE
ZIAGEN, ABACAVIR SULFATE

VINTAGE

* VINTAGE PHARMACEUTICALS LLC
ACETAMINOPHEN AND CODEINE PHOSPHATE, ACETAMINOPHEN
BENZTROPINE MESYLATE, BENZTROPINE MESYLATE
FOLIC ACID, FOLIC ACID
HYDROCORTISONE AND ACETIC ACID, ACETIC ACID, GLACIAL
HYDROCORTISONE, HYDROCORTISONE
NYSTATIN, NYSTATIN
PHENYLEPHRINE HYDROCHLORIDE AND PROMETHAZINE HYDROCHLORIDE, PHENYLEPHRINE
PREDNISOLONE SODIUM PHOSPHATE, PREDNISOLONE SODIUM PHOSPHATE
PROMETH HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE W/CODEINE PHOSPHATE, CODEINE
PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
PROMETHAZINE WITH CODEINE, CODEINE PHOSPHATE
ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE

VINTAGE PHARMS

* VINTAGE PHARMACEUTICALS
ALPRAZOLAM, ALPRAZOLAM
CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
CYCLAFEM 0.5/35, ETHINYL ESTRADIOL
DEXTROAMPHETAMINE SULFATE, DEXTROAMPHETAMINE SULFATE
GILDAGIA, ETHINYL ESTRADIOL
GILDESS 24 FE, ETHINYL ESTRADIOL
GRISEOFULVIN, GRISEOFULVIN, MICROSIZE
HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
KIMIDESS, DESOGESTREL
PROPRANOLOL HYDROCHLORIDE, PROPRANOLOL HYDROCHLORIDE

* VINTAGE PHARMACEUTICALS INC
ACETAMINOPHEN AND CODEINE PHOSPHATE, ACETAMINOPHEN
ALLOPURINOL, ALLOPURINOL
AMITRIPTYLINE HYDROCHLORIDE, AMITRIPTYLINE HYDROCHLORIDE
BACLOFEN, BACLOFEN
BUTALBITAL, ACETAMINOPHEN AND CAFFEINE, ACETAMINOPHEN
BUTALBITAL, ACETAMINOPHEN, CAFFEINE AND CODEINE PHOSPHATE, ACETAMINOPHEN
DIAZEPAM, DIAZEPAM
HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
HYDROCODONE BITARTRATE AND IBUPROFEN, HYDROCODONE BITARTRATE
HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE
IBUPROFEN, IBUPROFEN
IBUPROFEN, IBUPROFEN (OTC)
ISOSORBIDE MONONITRATE, ISOSORBIDE MONONITRATE
LEVETIRACETAM, LEVETIRACETAM
MEPERIDINE HYDROCHLORIDE, MEPERIDINE HYDROCHLORIDE
METHYLPREDNISOLONE, METHYLPREDNISOLONE
METOCLOPRAMIDE HYDROCHLORIDE, METOCLOPRAMIDE HYDROCHLORIDE
OXYBUTYNYN CHLORIDE, OXYBUTYNYN CHLORIDE
OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN
PERPHENAZINE, PERPHENAZINE
PREDNISONE, PREDNISONE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** V ****

* VINTAGE PHARMACEUTICALS INC
 PROPAFENONE HYDROCHLORIDE, PROPAFENONE HYDROCHLORIDE
 SULFASALAZINE, SULFASALAZINE
 TORSEMIDE, TORSEMIDE

VINTAGE PHARMS LLC

* VINTAGE PHARMACEUTICALS LLC
 CYCLAFEM 1/35, ETHINYL ESTRADIOL
 CYCLAFEM 7/7/7, ETHINYL ESTRADIOL
 DUTASTERIDE, DUTASTERIDE
 EMOQUETTE, DESOGESTREL
 FELODIPINE, FELODIPINE
 GILDESS 1.5/30, ETHINYL ESTRADIOL
 GILDESS 1/20, ETHINYL ESTRADIOL
 GILDESS FE 1.5/30, ETHINYL ESTRADIOL
 GILDESS FE 1/20, ETHINYL ESTRADIOL
 LETROZOLE, LETROZOLE
 MORPHINE SULFATE, MORPHINE SULFATE
 MYCOPHENOLATE MOFETIL, MYCOPHENOLATE MOFETIL
 MYZILRA, ETHINYL ESTRADIOL
 ORSYTHIA, ETHINYL ESTRADIOL
 PERCOCET, ACETAMINOPHEN
 PREVIFEM, ETHINYL ESTRADIOL
 TRI-PREVIFEM, ETHINYL ESTRADIOL

VIOKACE

* VIOKACE LLC
 VIOKACE, PANCRELIPASE (AMYLASE)

VIRTUS PHARM

* VIRTUS PHARMACEUTICAL INC
 ACARBOSE, ACARBOSE
 ALBUTEROL SULFATE, ALBUTEROL SULFATE
 PALONOSETRON HYDROCHLORIDE, PALONOSETRON HYDROCHLORIDE

VIRTUS PHARMS

* VIRTUS PHARMACEUTICALS LLC
 DAPSONE, DAPSONE
 LEVORPHANOL TARTRATE, LEVORPHANOL TARTRATE
 PHENDIMETRAZINE TARTRATE, PHENDIMETRAZINE TARTRATE
 PROMETRIUM, PROGESTERONE

VISTA PHARMS

* VISTA PHARMACEUTICALS INC
 SULFAMETHOXAZOLE AND TRIMETHOPRIM, SULFAMETHOXAZOLE

VISTAPHARM

* VISTAPHARM INC
 ALBUTEROL SULFATE, ALBUTEROL SULFATE
 AMINOCAPROIC ACID, AMINOCAPROIC ACID
 ARIPIPIRAZOLE, ARIPIPIRAZOLE
 CHLORTHALIDONE, CHLORTHALIDONE
 CLOBAZAM, CLOBAZAM
 DIGOXIN, DIGOXIN
 FELBAMATE, FELBAMATE
 GABAPENTIN, GABAPENTIN
 HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
 LACTULOSE, LACTULOSE
 METHADONE HYDROCHLORIDE, METHADONE HYDROCHLORIDE
 METOCLOPRAMIDE HYDROCHLORIDE, METOCLOPRAMIDE HYDROCHLORIDE
 MYCOPHENOLATE MOFETIL, MYCOPHENOLATE MOFETIL
 NYSTATIN, NYSTATIN
 OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
 PHENYTOIN, PHENYTOIN
 PREDNISOLONE, PREDNISOLONE
 VALPROIC ACID, VALPROIC ACID

VITRUVIAS THERAP

* VITRUVIAS THERAPEUTICS
 POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
 * VITRUVIAS THERAPEUTICS LLC
 CYANOCOBALAMIN, CYANOCOBALAMIN

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** V ****

* VITRUVIAS THERAPEUTICS LLC
LIDOCAINE, LIDOCAINE
NYSTATIN AND TRIAMCINOLONE ACETONIDE, NYSTATIN

VIVUS

* VIVUS INC
QSYMIA, PHENTERMINE HYDROCHLORIDE

VIVUS INC

* VIVUS INC
PANCREAZE, PANCRELIPASE (AMYLASE)

VKT PHARMA PVT LTD

* VKT PHARMA PRIVATE LTD
RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE

VPNA

* VALEANT PHARMACEUTICALS NORTH AMERICA
DICLOFENAC SODIUM, DICLOFENAC SODIUM

VYERA PHARMS LLC

* VYERA PHARMACEUTICALS LLC
DARAPRIM, PYRIMETHAMINE

**** W ******WA UNIV SCH MED**

* WASHINGTON UNIV SCHOOL MEDICINE
AMMONIA N 13, AMMONIA N-13
CHOLINE C-11, CHOLINE C-11

WATSON LABS

* WATSON LABORATORIES
FOLIC ACID, FOLIC ACID
PROPAPENONE HYDROCHLORIDE, PROPAPENONE HYDROCHLORIDE

* WATSON LABORATORIES INC
ACARBOSE, ACARBOSE
ALBUTEROL SULFATE, ALBUTEROL SULFATE
ALENDRONATE SODIUM, ALENDRONATE SODIUM
ALLOPURINOL, ALLOPURINOL
AMLODIPINE BESYLATE AND BENAZEPRIL HYDROCHLORIDE, AMLODIPINE BESYLATE
AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
AMOXAPINE, AMOXAPINE
ATENOLOL AND CHLORTHALIDONE, ATENOLOL
CAPTOPRIL, CAPTOPRIL
CARISOPRODOL, CARISOPRODOL
CHLORZOXAZONE, CHLORZOXAZONE
CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE
CLINDAMYCIN HYDROCHLORIDE, CLINDAMYCIN HYDROCHLORIDE
CLONAZEPAM, CLONAZEPAM
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
HYDROXYCHLOROQUINE SULFATE, HYDROXYCHLOROQUINE SULFATE
IPRATROPIUM BROMIDE, IPRATROPIUM BROMIDE
LABETALOL HYDROCHLORIDE, LABETALOL HYDROCHLORIDE
LAMOTRIGINE, LAMOTRIGINE
LEVONORGESTREL AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
LISINAPRIL, LISINAPRIL
LORAZEPAM, LORAZEPAM
LOSARTAN POTASSIUM, LOSARTAN POTASSIUM
LOXAPINE SUCCINATE, LOXAPINE SUCCINATE
MEPROBAMATE, MEPROBAMATE
METHYLDOPA, METHYLDOPA
METHYLPREDNISOLONE, METHYLPREDNISOLONE
METRONIDAZOLE, METRONIDAZOLE
MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** W ****

- * WATSON LABORATORIES INC
 MINOXIDIL, MINOXIDIL
 NABUMETONE, NABUMETONE
 NALOXONE HYDROCHLORIDE AND PENTAZOCINE HYDROCHLORIDE, NALOXONE HYDROCHLORIDE
 NATEGLINIDE, NATEGLINIDE
 NIZATIDINE, NIZATIDINE
 NORETHINDRONE AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
 OGESTREL 0.5/50-28, ETHINYL ESTRADIOL
 ORPHENADRINE CITRATE, ORPHENADRINE CITRATE
 OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN
 PRAVASTATIN SODIUM, PRAVASTATIN SODIUM
 PREDNISOLONE, PREDNISOLONE
 PREDNISONE, PREDNISONE
 PRIMIDONE, PRIMIDONE
 PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
 PROPRANOLOL HYDROCHLORIDE, PROPRANOLOL HYDROCHLORIDE
 QUASENSE, ETHINYL ESTRADIOL
 QUINIDINE SULFATE, QUINIDINE SULFATE
 RAMIPRIL, RAMIPRIL
 RIVASTIGMINE TARTRATE, RIVASTIGMINE TARTRATE
 SULFASALAZINE, SULFASALAZINE
 SULINDAC, SULINDAC
 SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
 TETRACYCLINE HYDROCHLORIDE, TETRACYCLINE HYDROCHLORIDE
 TOPIRAMATE, TOPIRAMATE
 TRIAMTERENE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 TRIHEXYPHENIDYL HYDROCHLORIDE, TRIHEXYPHENIDYL HYDROCHLORIDE
 TRIMETHOPRIM, TRIMETHOPRIM
 VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE
 VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE
 ZOVIA 1/50E-28, ETHINYL ESTRADIOL
- * WATSON LABS INC
 LISINAPRIL AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE

WATSON LABS INC

- * WATSON LABORATORIES INC
 AMANTADINE HYDROCHLORIDE, AMANTADINE HYDROCHLORIDE
 AMBRISENTAN, AMBRISENTAN
 AMLODIPINE BESYLATE AND BENAZEPRIL HYDROCHLORIDE, AMLODIPINE BESYLATE
 AMMONIUM LACTATE, AMMONIUM LACTATE
 BOSENTAN, BOSENTAN
 BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
 CELECOXIB, CELECOXIB
 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
 MOXIFLOXACIN HYDROCHLORIDE, MOXIFLOXACIN HYDROCHLORIDE
 NITROFURANTOIN (MONOHYDRATE/MACROCRYSTALS), NITROFURANTOIN
 OLOPATADINE HYDROCHLORIDE, OLOPATADINE HYDROCHLORIDE
 PENICILLAMINE, PENICILLAMINE
 PERPHENAZINE, PERPHENAZINE
 PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
 PROPAFENONE HYDROCHLORIDE, PROPAFENONE HYDROCHLORIDE
 PROPOFOL, PROPOFOL
 RALOXIFENE HYDROCHLORIDE, RALOXIFENE HYDROCHLORIDE
 RASAGILINE MESYLATE, RASAGILINE MESYLATE
 ROPINIROLE HYDROCHLORIDE, ROPINIROLE HYDROCHLORIDE
 ROSUVASTATIN CALCIUM, ROSUVASTATIN CALCIUM
 SILDENAFIL CITRATE, SILDENAFIL CITRATE
 SOLIFENACIN SUCCINATE, SOLIFENACIN SUCCINATE

WATSON LABS TEVA

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** W ****

* WATSON LABORATORIES INC AN INDIRECT WHOLLY OWNED SUB OF TEVA PHARMACEUTICALS USA INC
 ALBUTEROL SULFATE AND IPRATROPIUM BROMIDE, ALBUTEROL SULFATE
 ALVIMOPAN, ALVIMOPAN
 BICALUTAMIDE, BICALUTAMIDE
 BUPRENORPHINE, BUPRENORPHINE
 CINACALCET HYDROCHLORIDE, CINACALCET HYDROCHLORIDE
 EZETIMIBE AND ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM
 GLIPIZIDE, GLIPIZIDE
 IBANDRONATE SODIUM, IBANDRONATE SODIUM
 ISRADIPINE, ISRADIPINE
 LEVOFLOXACIN, LEVOFLOXACIN
 MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE
 NORETHINDRONE AND ETHINYL ESTRADIOL (10/11), ETHINYL ESTRADIOL
 NORETHINDRONE AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
 PROBENECID, PROBENECID
 SIMVASTATIN, SIMVASTATIN
 TEMOZOLOMIDE, TEMOZOLOMIDE
 TRIENTINE HYDROCHLORIDE, TRIENTINE HYDROCHLORIDE

WATSON PHARMS INC

* WATSON PHARMACEUTICALS INC
 TESTOSTERONE CYPIONATE, TESTOSTERONE CYPIONATE
 TESTOSTERONE ENANTHATE, TESTOSTERONE ENANTHATE

WATSON PHARMS TEVA

* WATSON PHARMACEUTICALS INC AN INDIRECT WHOLLY OWNED SUB OF TEVA PHARMACEUTICALS USA INC
 RIFAMPIN, RIFAMPIN

WELLSTAT THERAP

* WELLSTAT THERAPEUTICS CORP
 VISTOGARD, URIDINE TRIACETATE
 XURIDEN, URIDINE TRIACETATE

WES PHARMA INC

* WES PHARMA INC
 HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
 OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN

WEST WARD

* WEST WARD PHARMACEUTICAL CORP
 DICYCLOMINE HYDROCHLORIDE, DICYCLOMINE HYDROCHLORIDE

WEST WARD PHARM CORP

* WEST WARD PHARMACEUTICAL CORP
 PHENYLEPHRINE HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE
 ROCURONIUM BROMIDE, ROCURONIUM BROMIDE

WEST-WARD PHARMS INT

* WEST-WARD PHARMACEUTICALS INTERNATIONAL LTD
 ACETAZOLAMIDE SODIUM, ACETAZOLAMIDE SODIUM
 ADENOSINE, ADENOSINE
 ALLOPURINOL SODIUM, ALLOPURINOL SODIUM
 ALPROSTADIL, ALPROSTADIL
 AMIKACIN SULFATE, AMIKACIN SULFATE
 AMPICILLIN AND SULBACTAM, AMPICILLIN SODIUM
 AMRINONE LACTATE, INAMRINONE LACTATE
 ATRACURIUM BESYLATE PRESERVATIVE FREE, ATRACURIUM BESYLATE
 ATRACURIUM BESYLATE, ATRACURIUM BESYLATE
 AZATHIOPRINE SODIUM, AZATHIOPRINE SODIUM
 BLEOMYCIN SULFATE, BLEOMYCIN SULFATE
 BUMETANIDE, BUMETANIDE
 BUPRENORPHINE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE
 BUTORPHANOL TARTRATE PRESERVATIVE FREE, BUTORPHANOL TARTRATE
 BUTORPHANOL TARTRATE, BUTORPHANOL TARTRATE
 CARBOPLATIN, CARBOPLATIN
 CEFOXITIN, CEFOXITIN SODIUM
 CERUBIDINE, DAUNORUBICIN HYDROCHLORIDE
 CHLOROPROCAINE HYDROCHLORIDE, CHLOROPROCAINE HYDROCHLORIDE
 CHLORPROMAZINE HYDROCHLORIDE, CHLORPROMAZINE HYDROCHLORIDE
 CISPLATIN, CISPLATIN
 CLADRIBINE, CLADRIBINE
 CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** W **

* WEST-WARD PHARMACEUTICALS INTERNATIONAL LTD
 CYANOCOBALAMIN, CYANOCOBALAMIN
 CYCLOSPORINE, CYCLOSPORINE
 CYTARABINE, CYTARABINE
 DACARBAZINE, DACARBAZINE
 DAUNORUBICIN HYDROCHLORIDE, DAUNORUBICIN HYDROCHLORIDE
 DEFEROXAMINE MESYLATE, DEFEROXAMINE MESYLATE
 DEXAMETHASONE SODIUM PHOSPHATE, DEXAMETHASONE SODIUM PHOSPHATE
 DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE
 DEXRAZOXANE HYDROCHLORIDE, DEXRAZOXANE HYDROCHLORIDE
 DICYCLOMINE HYDROCHLORIDE (PRESERVATIVE FREE), DICYCLOMINE HYDROCHLORIDE
 DIGOXIN, DIGOXIN
 DIHYDROERGOTAMINE MESYLATE, DIHYDROERGOTAMINE MESYLATE
 DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
 DIPHENHYDRAMINE HYDROCHLORIDE, DIPHENHYDRAMINE HYDROCHLORIDE
 DIPYRIDAMOLE, DIPYRIDAMOLE
 DOBUTAMINE HYDROCHLORIDE, DOBUTAMINE HYDROCHLORIDE
 DOXORUBICIN HYDROCHLORIDE, DOXORUBICIN HYDROCHLORIDE
 DOXYCYCLINE, DOXYCYCLINE HYCLATE
 EPIRUBICIN HYDROCHLORIDE, EPIRUBICIN HYDROCHLORIDE
 ESMOLOL HYDROCHLORIDE, ESMOLOL HYDROCHLORIDE
 ETOMIDATE, ETOMIDATE
 ETOPOSIDE, ETOPOSIDE
 FAMOTIDINE PRESERVATIVE FREE, FAMOTIDINE
 FAMOTIDINE, FAMOTIDINE
 FENOLDOPAM MESYLATE, FENOLDOPAM MESYLATE
 FLOXURIDINE, FLOXURIDINE
 FLUCONAZOLE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, FLUCONAZOLE
 FLUCONAZOLE IN SODIUM CHLORIDE 0.9%, FLUCONAZOLE
 FLUMAZENIL, FLUMAZENIL
 FLUPHENAZINE DECANOATE, FLUPHENAZINE DECANOATE
 FOSPHENYTOIN SODIUM, FOSPHENYTOIN SODIUM
 GRANISETRON HYDROCHLORIDE, GRANISETRON HYDROCHLORIDE
 HALOPERIDOL DECANOATE, HALOPERIDOL DECANOATE
 HALOPERIDOL, HALOPERIDOL LACTATE
 IDARUBICIN HYDROCHLORIDE, IDARUBICIN HYDROCHLORIDE
 IFOSFAMIDE, IFOSFAMIDE
 INDOMETHACIN SODIUM, INDOMETHACIN SODIUM
 IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE
 KETAMINE HYDROCHLORIDE, KETAMINE HYDROCHLORIDE
 KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
 LABETALOL HYDROCHLORIDE, LABETALOL HYDROCHLORIDE
 LEUCOVORIN CALCIUM PRESERVATIVE FREE, LEUCOVORIN CALCIUM
 LEUCOVORIN CALCIUM, LEUCOVORIN CALCIUM
 LEVOCARNITINE, LEVOCARNITINE
 LEVOLEUCOVORIN CALCIUM, LEVOLEUCOVORIN CALCIUM
 MELPHALAN HYDROCHLORIDE, MELPHALAN HYDROCHLORIDE
 MEPERIDINE HYDROCHLORIDE PRESERVATIVE FREE, MEPERIDINE HYDROCHLORIDE
 MEPERIDINE HYDROCHLORIDE, MEPERIDINE HYDROCHLORIDE
 MESNA, MESNA
 METHOTREXATE SODIUM PRESERVATIVE FREE, METHOTREXATE SODIUM
 METHOTREXATE SODIUM, METHOTREXATE SODIUM
 METOPROLOL TARTRATE, METOPROLOL TARTRATE
 MIDAZOLAM HYDROCHLORIDE, MIDAZOLAM HYDROCHLORIDE
 MILRINONE LACTATE IN DEXTROSE 5% IN PLASTIC CONTAINER, MILRINONE LACTATE
 MILRINONE LACTATE, MILRINONE LACTATE
 MITOMYCIN, MITOMYCIN
 MITOXANTRONE HYDROCHLORIDE, MITOXANTRONE HYDROCHLORIDE
 NALOXONE, NALOXONE HYDROCHLORIDE
 NOREPINEPHRINE BITARTRATE, NOREPINEPHRINE BITARTRATE
 OCTREOTIDE ACETATE (PRESERVATIVE FREE), OCTREOTIDE ACETATE
 OCTREOTIDE ACETATE, OCTREOTIDE ACETATE
 ONDANSETRON HYDROCHLORIDE PRESERVATIVE FREE, ONDANSETRON HYDROCHLORIDE
 ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
 ORPHENADRINE CITRATE, ORPHENADRINE CITRATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** W ****

* WEST-WARD PHARMACEUTICALS INTERNATIONAL LTD
 OXYTOCIN, OXYTOCIN
 PACLITAXEL, PACLITAXEL
 PAMIDRONATE DISODIUM, PAMIDRONATE DISODIUM
 PENTOSTATIN, PENTOSTATIN
 PHENTOLAMINE MESYLATE, PHENTOLAMINE MESYLATE
 PHENYTOIN SODIUM, PHENYTOIN SODIUM
 PROCHLORPERAZINE EDISYLATE, PROCHLORPERAZINE EDISYLATE
 PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
 RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
 RIFAMPIN, RIFAMPIN
 SODIUM CHLORIDE 0.9%, SODIUM CHLORIDE
 SODIUM FERRIC GLUCONATE COMPLEX IN SUCROSE, SODIUM FERRIC GLUCONATE COMPLEX
 STERILE WATER FOR INJECTION, STERILE WATER FOR INJECTION
 SUFENTANIL CITRATE, SUFENTANIL CITRATE
 SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
 TESTOSTERONE CYPIONATE, TESTOSTERONE CYPIONATE
 THIOTEPA, THIOTEPA
 TOBRAMYCIN SULFATE, TOBRAMYCIN SULFATE
 VECURONIUM BROMIDE, VECURONIUM BROMIDE
 VINBLASTINE SULFATE, VINBLASTINE SULFATE
 VINOURELBINE TARTRATE, VINOURELBINE TARTRATE

WI MEDCL CYCLOTRON

* WISCONSIN MEDICAL CYCLOTRON LLC
 AMMONIA N 13, AMMONIA N-13
 FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18

WILSHIRE PHARMS INC

* WILSHIRE PHARMACEUTICALS INC
 CARISOPRODOL, CARISOPRODOL
 NATEGLINIDE, NATEGLINIDE
 PERPHENAZINE, PERPHENAZINE
 PROPAFENONE HYDROCHLORIDE, PROPAFENONE HYDROCHLORIDE
 SEVELAMER CARBONATE, SEVELAMER CARBONATE
 TESTOSTERONE CYPIONATE, TESTOSTERONE CYPIONATE
 TIAGABINE HYDROCHLORIDE, TIAGABINE HYDROCHLORIDE

WINDLAS HLTHCARE

* WINDLAS HEALTHCARE PVT LTD
 AMILORIDE HYDROCHLORIDE, AMILORIDE HYDROCHLORIDE

WOCKHARDT

* WOCKHARDT LTD
 AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE
 AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
 AZITHROMYCIN, AZITHROMYCIN
 BETAXOLOL HYDROCHLORIDE, BETAXOLOL HYDROCHLORIDE
 BETHANECHOL CHLORIDE, BETHANECHOL CHLORIDE
 CEFPROZIL, CEFPROZIL
 CEFTAZIDIME, CEFTAZIDIME
 CEFTRIAXONE, CEFTRIAXONE SODIUM
 CEFUROXIME AXETIL, CEFUROXIME AXETIL
 CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 CLARITHROMYCIN, CLARITHROMYCIN
 DIVALPROEX SODIUM, DIVALPROEX SODIUM
 FAMOTIDINE, FAMOTIDINE (OTC)
 FOSPHENYTOIN SODIUM, FOSPHENYTOIN SODIUM
 FUROSEMIDE, FUROSEMIDE
 KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
 LEVETIRACETAM, LEVETIRACETAM
 LISINAPRIL, LISINAPRIL
 METOPROLOL SUCCINATE, METOPROLOL SUCCINATE
 NIACIN, NIACIN
 ONDANSETRON HYDROCHLORIDE PRESERVATIVE FREE, ONDANSETRON HYDROCHLORIDE
 ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
 PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
 RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE (OTC)
 SERTRALINE HYDROCHLORIDE, SERTRALINE HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** W ******* WOCKHARDT LTD**

SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
TAMSULOSIN HYDROCHLORIDE, TAMSULOSIN HYDROCHLORIDE
TIMOLOL MALEATE, TIMOLOL MALEATE
VALACYCLOVIR HYDROCHLORIDE, VALACYCLOVIR HYDROCHLORIDE
VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE

WOCKHARDT BIO AG*** WOCKHARDT BIO AG**

ABIRATERONE ACETATE, ABIRATERONE ACETATE
ACETAMINOPHEN AND CODEINE PHOSPHATE, ACETAMINOPHEN
ACETIC ACID, ACETIC ACID, GLACIAL
AMANTADINE HYDROCHLORIDE, AMANTADINE HYDROCHLORIDE
AMOXICILLIN AND CLAVULANATE POTASSIUM, AMOXICILLIN
AMOXICILLIN, AMOXICILLIN
BROMFED-DM, BROMPHENIRAMINE MALEATE
CARBAMAZEPINE, CARBAMAZEPINE
CHLORHEXIDINE GLUCONATE, CHLORHEXIDINE GLUCONATE
CIMETIDINE HYDROCHLORIDE, CIMETIDINE HYDROCHLORIDE
CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
CROMOLYN SODIUM, CROMOLYN SODIUM
CYCLOSPORINE, CYCLOSPORINE
DECITABINE, DECITABINE
DEXAMETHASONE, DEXAMETHASONE
DEXCHLORPHENIRAMINE MALEATE, DEXCHLORPHENIRAMINE MALEATE
DOXEPIN HYDROCHLORIDE, DOXEPIN HYDROCHLORIDE
ERYTHROMYCIN, ERYTHROMYCIN
FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
FLUTICASONE PROPIONATE, FLUTICASONE PROPIONATE
FUROSEMIDE, FUROSEMIDE
GENERLAC, LACTULOSE
HYDROCODONE BITARTRATE AND HOMATROPINE METHYLBROMIDE, HOMATROPINE METHYLBROMIDE
HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE
IMATINIB MESYLATE, IMATINIB MESYLATE
LACTULOSE, LACTULOSE
LEVETIRACETAM, LEVETIRACETAM
LIDOCAINE HYDROCHLORIDE, LIDOCAINE HYDROCHLORIDE
LINDANE, LINDANE
LITHIUM CITRATE, LITHIUM CITRATE
LOPERAMIDE HYDROCHLORIDE, LOPERAMIDE HYDROCHLORIDE (OTC)
LORATADINE, LORATADINE (OTC)
MEGESTROL ACETATE, MEGESTROL ACETATE
METOCLOPRAMIDE HYDROCHLORIDE, METOCLOPRAMIDE HYDROCHLORIDE
MINOXIDIL (FOR MEN), MINOXIDIL (OTC)
MINOXIDIL EXTRA STRENGTH (FOR MEN), MINOXIDIL (OTC)
NYSTATIN, NYSTATIN
OXACILLIN SODIUM, OXACILLIN SODIUM
OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE
OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
PHENYTOIN, PHENYTOIN
PIPERACILLIN AND TAZOBACTAM, PIPERACILLIN SODIUM
PREDNISOLONE SODIUM PHOSPHATE, PREDNISOLONE SODIUM PHOSPHATE
PREDNISOLONE, PREDNISOLONE
PROMETHAZINE HYDROCHLORIDE AND CODEINE PHOSPHATE, CODEINE PHOSPHATE
PROMETHAZINE PLAIN, PROMETHAZINE HYDROCHLORIDE
PROMETHAZINE W/ DEXTROMETHORPHAN, DEXTROMETHORPHAN HYDROBROMIDE
SELENIUM SULFIDE, SELENIUM SULFIDE
TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE
VALPROIC ACID, VALPROIC ACID

WOCKHARDT LTD*** WOCKHARDT LTD**

BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
CAPTOPRIL, CAPTOPRIL
CHILDREN'S FEXOFENADINE HYDROCHLORIDE ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
CHILDREN'S FEXOFENADINE HYDROCHLORIDE HIVES, FEXOFENADINE HYDROCHLORIDE (OTC)
ENALAPRIL MALEATE, ENALAPRIL MALEATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** W ****

* WOCKHARDT LTD
 ENTACAPONE, ENTACAPONE
 FEXOFENADINE HYDROCHLORIDE ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
 FEXOFENADINE HYDROCHLORIDE HIVES, FEXOFENADINE HYDROCHLORIDE (OTC)
 LAMOTRIGINE, LAMOTRIGINE
 LANSOPRAZOLE, LANSOPRAZOLE (OTC)
 OLOPATADINE HYDROCHLORIDE, OLOPATADINE HYDROCHLORIDE
 RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
 ZIPRASIDONE HYDROCHLORIDE, ZIPRASIDONE HYDROCHLORIDE

WOCKHARDT USA

* WOCKHARDT USA INC
 GRANISETRON HYDROCHLORIDE, GRANISETRON HYDROCHLORIDE
 * WOCKHARDT USA LLC
 LANSOPRAZOLE, LANSOPRAZOLE

WOODWARD

* WOODWARD PHARMA SERVICES LLC
 FLUCONAZOLE IN DEXTROSE 5% IN PLASTIC CONTAINER, FLUCONAZOLE
 FLUCONAZOLE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, FLUCONAZOLE
 MILRINONE LACTATE IN DEXTROSE 5%, MILRINONE LACTATE

WRASER PHARMS

* WRASER PHARMACEUTICALS LLC
 CETRAXAL, CIPROFLOXACIN HYDROCHLORIDE

WRASER PHARMS LLC

* WRASER PHARMACEUTICALS LLC
 BUTALBITAL, ACETAMINOPHEN AND CAFFEINE, ACETAMINOPHEN
 TREZIX, ACETAMINOPHEN

WUSM CYCLOTRON

* WASHINGTON UNIV SCH MEDICINE CYCLOTRON FACILITY
 FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18

WYETH PHARMS

* WYETH PHARMACEUTICALS LLC
 DUAVEE, BAZEDOXIFENE ACETATE
 EFFEXOR XR, VENLAFAXINE HYDROCHLORIDE
 PHOSPHOLINE IODIDE, ECHOTHIOPHATE IODIDE
 PREMARIN, ESTROGENS, CONJUGATED
 PREMPHASE 14/14, ESTROGENS, CONJUGATED
 PREMPRO, ESTROGENS, CONJUGATED
 PROTONIX IV, PANTOPRAZOLE SODIUM
 PROTONIX, PANTOPRAZOLE SODIUM
 TRECATOR, ETHIONAMIDE
 ZOSYN IN PLASTIC CONTAINER, PIPERACILLIN SODIUM
 ZOSYN, PIPERACILLIN SODIUM

**** X ******X GEN PHARMS**

* X GEN PHARMACEUTICALS INC
 BACIIM, BACITRACIN

XELLIA PHARMS APS

* XELLIA PHARMACEUTICALS APS
 BACITRACIN, BACITRACIN
 CASPOFUNGIN ACETATE, CASPOFUNGIN ACETATE
 COLISTIMETHATE SODIUM, COLISTIMETHATE SODIUM
 DAPTOMYCIN, DAPTOMYCIN
 POLYMYXIN B SULFATE, POLYMYXIN B SULFATE
 TIGECYCLINE, TIGECYCLINE
 TOBRAMYCIN SULFATE, TOBRAMYCIN SULFATE
 VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE
 VORICONAZOLE, VORICONAZOLE

XERIS

* XERIS PHARMACEUTICALS INC
 GVOKE HYPOPEN, GLUCAGON
 GVOKE PFS, GLUCAGON

XGEN PHARMS

* XGEN PHARMACEUTICALS DJB INC

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** X ****

* XGEN PHARMACEUTICALS DJB INC
 ACETAZOLAMIDE SODIUM, ACETAZOLAMIDE SODIUM
 AMPHOTERICIN B, AMPHOTERICIN B
 CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
 COLISTIMETHATE SODIUM, COLISTIMETHATE SODIUM
 DACTINOMYCIN, DACTINOMYCIN
 FOLIC ACID, FOLIC ACID
 HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
 IBUPROFEN LYSINE, IBUPROFEN LYSINE
 LEVETIRACETAM, LEVETIRACETAM
 LINCOMYCIN, LINCOMYCIN HYDROCHLORIDE
 LIOTHYRONINE SODIUM, LIOTHYRONINE SODIUM
 NEOMYCIN AND POLYMYXIN B SULFATE, NEOMYCIN SULFATE
 NEOMYCIN SULFATE, NEOMYCIN SULFATE
 NYSTATIN, NYSTATIN
 POLYMYXIN B SULFATE, POLYMYXIN B SULFATE
 PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
 STREPTOMYCIN SULFATE, STREPTOMYCIN SULFATE
 TOBRAMYCIN SULFATE, TOBRAMYCIN SULFATE
 TRANEXAMIC ACID, TRANEXAMIC ACID

XIAMEN LP PHARM CO

* XIAMEN LP PHARMACUETICAL CO LTD
 CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE

XIROMED

* XIROMED PHARMA ESPANA SL
 ACYCLOVIR, ACYCLOVIR
 ALTAVERA, ETHINYL ESTRADIOL
 CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
 ELIFEMME, ETHINYL ESTRADIOL
 ESTARYLLA, ETHINYL ESTRADIOL
 FLUOCINONIDE, FLUOCINONIDE
 HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
 INTROVALE, ETHINYL ESTRADIOL
 ISIBLOOM, DESOGESTREL
 JAIMIESS, ETHINYL ESTRADIOL
 LANSOPRAZOLE, LANSOPRAZOLE
 LEVONORGESTREL AND ETHINYL ESTRADIOL AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
 LEVONORGESTREL, LEVONORGESTREL (OTC)
 LORYNA, DROSPIRENONE
 NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL AND FERROUS FUMARATE, ETHINYL ESTRADIOL
 OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN
 PROGESTERONE, PROGESTERONE
 SYEDA, DROSPIRENONE
 TRI-ESTARYLLA, ETHINYL ESTRADIOL
 TRI-LO-ESTARYLLA, ETHINYL ESTRADIOL
 VIENVA, ETHINYL ESTRADIOL
 VOLNEA, DESOGESTREL

XSPIRE PHARMA

* XSPIRE PHARMA
 NALFON, FENOPROFEN CALCIUM
 * XSPIRE PHARMA LLC
 DEXAMETHASONE, DEXAMETHASONE
 FENOPROFEN CALCIUM, FENOPROFEN CALCIUM

XTTRIUM

* XTTRIUM LABORATORIES INC
 CHLORHEXIDINE GLUCONATE, CHLORHEXIDINE GLUCONATE
 EXIDINE, CHLORHEXIDINE GLUCONATE (OTC)

XTTRIUM LABS INC

* XTTRIUM LABORATORIES INC
 LACTULOSE, LACTULOSE

**** Y ******YABAO PHARM**

* YABAO PHARMACEUTICAL CO LTD BEIJING
 GALANTAMINE HYDROBROMIDE, GALANTAMINE HYDROBROMIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** Y ******YAOPHARMA CO LTD**

- * YAOPHARMA CO LTD
 - DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE
 - ENTECAVIR, ENTECAVIR
 - VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE

YICHANG HUMANWELL

- * YICHANG HUMANWELL PHARMACEUTICAL CO LTD
 - BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
 - NAPROXEN SODIUM, NAPROXEN SODIUM (OTC)
 - POTASSIUM CHLORIDE, POTASSIUM CHLORIDE

YILING PHARM LTD

- * YILING PHARMACEUTICAL LTD
 - ACYCLOVIR, ACYCLOVIR
 - BUSPIRONE HYDROCHLORIDE, BUSPIRONE HYDROCHLORIDE
 - CELECOXIB, CELECOXIB
 - CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE
 - FELODIPINE, FELODIPINE
 - VALACYCLOVIR HYDROCHLORIDE, VALACYCLOVIR HYDROCHLORIDE

YOUNGTECH PHARMS INC

- * YOUNGTECH PHARMACEUTICALS INC
 - METOPROLOL TARTRATE, METOPROLOL TARTRATE

YUNG SHIN PHARM

- * YUNG SHIN PHARMACEUTICAL INDUSTRIAL CO LTD
 - CEFACTOR, CEFACTOR
 - CEPHALEXIN, CEPHALEXIN
 - CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
 - DICLOFENAC SODIUM AND MISOPROSTOL, DICLOFENAC SODIUM
 - FELODIPINE, FELODIPINE
 - GRANISETRON HYDROCHLORIDE, GRANISETRON HYDROCHLORIDE
 - MELOXICAM, MELOXICAM
 - ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE

**** Z ******ZAMBON SPA**

- * ZAMBON SPA ITALY
 - MONUROL, FOSFOMYCIN TROMETHAMINE

ZENNOVA

- * ZENNOVA LLC
 - IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE
- * ZENNOVA PHARMACEUTICALS CHENGDU CO LTD
 - PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE
 - QUETIAPINE FUMARATE, QUETIAPINE FUMARATE

ZENPEP

- * ZENPEP LLC
 - ZENPEP, PANCRELIPASE (AMYLASE)

ZHEJIANG HISUN PHARM

- * ZHEJIANG HISUN PHARMACEUTICAL CO LTD
 - MYCOPHENOLATE MOFETIL, MYCOPHENOLATE MOFETIL

ZHEJIANG JUTAI PHARM

- * ZHEJIANG JUTAI PHARMACEUTICAL CO LTD
 - BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE

ZHEJIANG YONGTAI

- * ZHEJIANG YONGTAI PHARMACEUTICAL CO LTD
 - ROSUVASTATIN CALCIUM, ROSUVASTATIN CALCIUM

ZO SKIN HEALTH

- * ZO SKIN HEALTH
 - TRETINOIN, TRETINOIN

ZYDUS

- * ZYDUS WORLDWIDE DMCC
 - AZITHROMYCIN, AZITHROMYCIN
 - BACLOFEN, BACLOFEN
 - CHLORPROMAZINE HYDROCHLORIDE, CHLORPROMAZINE HYDROCHLORIDE
 - ISOSORBIDE DINITRATE, ISOSORBIDE DINITRATE
 - LEFLUNOMIDE, LEFLUNOMIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** Z ****

* ZYDUS WORLDWIDE DMCC

MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE
 PHYTONADIONE, PHYTONADIONE
 URSODIOL, URSODIOL

ZYDUS HLTHCARE

* ZYDUS HEALTHCARE USA LLC

DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE
 LANSOPRAZOLE, LANSOPRAZOLE
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE

ZYDUS NOVELTECH INC

* ZYDUS NOVELTECH INC

RIVASTIGMINE, RIVASTIGMINE

ZYDUS PHARMS

* ZYDUS PHARMACEUTICALS USA INC

ACAMPROSATE CALCIUM, ACAMPROSATE CALCIUM
 ACETAZOLAMIDE SODIUM, ACETAZOLAMIDE SODIUM
 ACETYLCYSTEINE, ACETYLCYSTEINE
 ACYCLOVIR SODIUM, ACYCLOVIR SODIUM
 ACYCLOVIR, ACYCLOVIR
 ALBENDAZOLE, ALBENDAZOLE
 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
 ARIPIPIRAZOLE, ARIPIPIRAZOLE
 ARSENIC TRIOXIDE, ARSENIC TRIOXIDE
 ASPIRIN AND DIPYRIDAMOLE, ASPIRIN
 ATENOLOL AND CHLORTHALIDONE, ATENOLOL
 ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM
 AZELASTINE HYDROCHLORIDE, AZELASTINE HYDROCHLORIDE
 BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
 BOSENTAN, BOSENTAN
 BUDESONIDE, BUDESONIDE
 BUMETANIDE, BUMETANIDE
 BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
 BUSPIRONE HYDROCHLORIDE, BUSPIRONE HYDROCHLORIDE
 CANDESARTAN CILEXETIL AND HYDROCHLOROTHIAZIDE, CANDESARTAN CILEXETIL
 CANDESARTAN CILEXETIL, CANDESARTAN CILEXETIL
 CARBAMAZEPINE, CARBAMAZEPINE
 CARBIDOPA, CARBIDOPA
 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
 COLESEVELAM HYDROCHLORIDE, COLESEVELAM HYDROCHLORIDE
 CYPROHEPTADINE HYDROCHLORIDE, CYPROHEPTADINE HYDROCHLORIDE
 DEFERASIROX, DEFERASIROX
 DESMOPRESSIN ACETATE (NEEDS NO REFRIGERATION), DESMOPRESSIN ACETATE
 DESOXIMETASONE, DESOXIMETASONE
 DESVENLAFAXINE SUCCINATE, DESVENLAFAXINE SUCCINATE
 DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE
 DIFLUNISAL, DIFLUNISAL
 DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
 DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
 DOXAZOSIN MESYLATE, DOXAZOSIN MESYLATE
 DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
 DOXYCYCLINE, DOXYCYCLINE
 DOXYCYCLINE, DOXYCYCLINE HYCLATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** Z **

* ZYDUS PHARMACEUTICALS USA INC
 DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE
 DUTASTERIDE AND TAMSULOSIN HYDROCHLORIDE, DUTASTERIDE
 ELETRIPTAN HYDROBROMIDE, ELETRIPTAN HYDROBROMIDE
 EMTRICITABINE AND TENOFOVIR DISOPROXIL FUMARATE, EMTRICITABINE
 ENTECAVIR, ENTECAVIR
 ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM
 ETHACRYNATE SODIUM, ETHACRYNATE SODIUM
 ETODOLAC, ETODOLAC
 ETOMIDATE, ETOMIDATE
 EXEMESTANE, EXEMESTANE
 EZETIMIBE, EZETIMIBE
 FESOTERODINE FUMARATE, FESOTERODINE FUMARATE
 FLUCONAZOLE, FLUCONAZOLE
 FLUOCINONIDE, FLUOCINONIDE
 GLIPIZIDE, GLIPIZIDE
 GLYBURIDE AND METFORMIN HYDROCHLORIDE, GLYBURIDE
 GLYBURIDE, GLYBURIDE
 HALOPERIDOL DECANOATE, HALOPERIDOL DECANOATE
 INDOMETHACIN, INDOMETHACIN
 ITRACONAZOLE, ITRACONAZOLE
 LABETALOL HYDROCHLORIDE, LABETALOL HYDROCHLORIDE
 LANSOPRAZOLE, LANSOPRAZOLE
 LINEZOLID, LINEZOLID
 MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
 MESALAMINE, MESALAMINE
 METHOTREXATE SODIUM, METHOTREXATE SODIUM
 METHYLPREDNISOLONE, METHYLPREDNISOLONE
 METOPROLOL SUCCINATE, METOPROLOL SUCCINATE
 MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE
 MIRTAZAPINE, MIRTAZAPINE
 MYCOPHENOLATE MOFETIL HYDROCHLORIDE, MYCOPHENOLATE MOFETIL HYDROCHLORIDE
 NADOLOL, NADOLOL
 NATEGLINIDE, NATEGLINIDE
 NIFEDIPINE, NIFEDIPINE
 NITROFURANTOIN, NITROFURANTOIN, MACROCRYSTALLINE
 OLANZAPINE, OLANZAPINE
 OLMESARTAN MEDOXOMIL, OLMESARTAN MEDOXOMIL
 OMEPRAZOLE AND SODIUM BICARBONATE, OMEPRAZOLE
 OMEPRAZOLE AND SODIUM BICARBONATE, OMEPRAZOLE (OTC)
 OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE
 PHENTERMINE HYDROCHLORIDE, PHENTERMINE HYDROCHLORIDE
 POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
 POTASSIUM CITRATE, POTASSIUM CITRATE
 PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE
 PYRIDOSTIGMINE BROMIDE, PYRIDOSTIGMINE BROMIDE
 RAMELTEON, RAMELTEON
 SIROLIMUS, SIROLIMUS
 SPIRONOLACTONE, SPIRONOLACTONE
 SUCCINYLCHOLINE CHLORIDE, SUCCINYLCHOLINE CHLORIDE
 TADALAFIL, TADALAFIL
 TAMOXIFEN CITRATE, TAMOXIFEN CITRATE
 TELMISARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 TELMISARTAN, TELMISARTAN
 TEMOZOLOMIDE, TEMOZOLOMIDE
 TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE
 TOPIRAMATE, TOPIRAMATE
 TRANEXAMIC ACID, TRANEXAMIC ACID
 TRAZODONE HYDROCHLORIDE, TRAZODONE HYDROCHLORIDE
 TRIAMTERENE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 TRIENTINE HYDROCHLORIDE, TRIENTINE HYDROCHLORIDE
 VALACYCLOVIR HYDROCHLORIDE, VALACYCLOVIR HYDROCHLORIDE
 VARDENAFIL HYDROCHLORIDE, VARDENAFIL HYDROCHLORIDE
 VORICONAZOLE, VORICONAZOLE
 ZOLMITRIPTAN, ZOLMITRIPTAN

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** Z ******ZYDUS PHARMS USA**

* ZYDUS PHARMACEUTICALS USA INC
 AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
 ATENOLOL, ATENOLOL
 AZATHIOPRINE, AZATHIOPRINE
 BENAZEPRIL HYDROCHLORIDE, BENAZEPRIL HYDROCHLORIDE
 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
 ZONISAMIDE, ZONISAMIDE

ZYDUS PHARMS USA INC

* ZYDUS PHARMACEUTICALS USA INC
 AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE
 ANASTROZOLE, ANASTROZOLE
 ATOMOXETINE HYDROCHLORIDE, ATOMOXETINE HYDROCHLORIDE
 BICALUTAMIDE, BICALUTAMIDE
 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
 LEVOFLOXACIN, LEVOFLOXACIN
 LOSARTAN POTASSIUM AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 LOSARTAN POTASSIUM, LOSARTAN POTASSIUM
 OMEPRAZOLE, OMEPRAZOLE
 PIOGLITAZONE HYDROCHLORIDE, PIOGLITAZONE HYDROCHLORIDE
 PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE
 PROPRANOLOL HYDROCHLORIDE, PROPRANOLOL HYDROCHLORIDE
 RANITIDINE HYDROCHLORIDE, RANITIDINE 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
 OXAYDO, OXYCODONE HYDROCHLORIDE
 SPRIX, KETOROLAC TROMETHAMINE
 VIVLODEX, MELOXICAM
 ZORVOLEX, DICLOFENAC

APPENDIX C**UNIFORM TERMS*****DOSAGE FORMS***

AEROSOL, FOAM	OINTMENT
AEROSOL, METERED	OINTMENT, AUGMENTED
BAR, CHEWABLE	PASTE
CAPSULE	PATCH
CAPSULE, DELAYED REL PELLETS	PELLET
CAPSULE, DELAYED RELEASE	PELLETS
CAPSULE, EXTENDED RELEASE	POWDER
CLOTH	POWDER, EXTENDED RELEASE
CONCENTRATE	POWDER, METERED
CREAM	RING
CREAM, AUGMENTED	SHAMPOO
ELIXIR	SOLUTION
EMULSION	SOLUTION FOR SLUSH
ENEMA	SOLUTION, EXTENDED RELEASE
FILM	SOLUTION, GEL FORMING/DROPS
FILM, EXTENDED RELEASE	SOLUTION, METERED
FOAM	SOLUTION/DROPS
FOR SOLUTION	SPONGE
FOR SUSPENSION	SPRAY
FOR SUSPENSION, DELAYED RELEASE	SPRAY, METERED
FOR SUSPENSION, EXTENDED RELEASE	SUPPOSITORY
GAS	SUSPENSION
GEL	SUSPENSION, EXTENDED RELEASE
GEL, AUGMENTED	SUSPENSION, LIPOSOMAL
GEL, METERED	SUSPENSION/DROPS
GRANULE	SWAB
GRANULE, DELAYED RELEASE	SYRUP
GUM, CHEWING	SYSTEM
IMPLANT	SYSTEM, EXTENDED RELEASE
INHALANT	TABLET
INJECTABLE	TABLET, CHEWABLE
INJECTABLE, LIPID COMPLEX	TABLET, DELAYED RELEASE
INJECTABLE, LIPOSOMAL	TABLET, EFFERVESCENT
INJECTION, EXTENDED RELEASE	TABLET, EXTENDED RELEASE
INSERT	TABLET, EXTENDED RELEASE, CHEWABLE
INSERT, EXTENDED RELEASE	TABLET, FOR SUSPENSION
INTRAUTERINE DEVICE	TABLET, ORALLY DISINTEGRATING
JELLY	TABLET, ORALLY DISINTEGRATING, DELAYED RELEASE
LIQUID	TABLET, ORALLY DISINTEGRATING, EXTENDED RELEASE
LOTION	TAPE
LOTION, AUGMENTED	TROCHE/LOZENGE
LOTION/SHAMPOO	
OIL	
OIL/DROPS	

Note: Terms comprise currently marketed products

APPENDIX C**UNIFORM TERMS*****ROUTES OF ADMINISTRATION***

BUCCAL	INTRAVESICAL
DENTAL	INTRAVITREAL
ENDOCERVICAL	IRRIGATION
ENDOTRACHEAL	IV (INFUSION)
ENTERAL	N/A
IMPLANTATION	NASAL
INHALATION	OPHTHALMIC
INJECTION	ORAL
INTERSTITIAL	ORAL-21
INTRA-ANAL	ORAL-28
INTRA-ARTERIAL	OTIC
INTRA-ARTICULAR	PERFUSION, CARDIAC
INTRACRANIAL	PERIODONTAL
INTRADERMAL	RECTAL
INTRAMUSCULAR	SPINAL
INTRAOCULAR	SUBCUTANEOUS
INTRAOSSEOUS	SUBLINGUAL
INTRAPERITONEAL	TOPICAL
INTRAPLEURAL	TRANSDERMAL
INTRATHECAL	TRANSMUCOSAL
INTRATRACHEAL	URETHRAL
INTRAUTERINE	VAGINAL
INTRAVENOUS	

Note: Terms comprise currently marketed products

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
SQ CM	SQUARE CENTIMETER
SC	SUBCUTANEOUS
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 (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 five-year and three-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 Antibiotics 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 established name of the active ingredient, followed by the proprietary name (brand name or trade name) of the drug product. Active ingredient headings for multiple 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.

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 patent information be filed with all newly submitted Section 505(b) drug applications. In addition, patent information must be filed on Form FDA 3542 within 30 days of the date of approval of a Section 505(b) drug application.¹ FDA publishes certain information from Form FDA 3542 in the Orange Book after approval of the NDA or supplement.

The Orange Book includes the patent submission date (i.e., the date on which the FDA receives patent information from the NDA holder) 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 ANDA.²

The patents that FDA regards as covered by the statutory provisions for submission of patent information are:

- Patents that claim the drug substance (i.e., active ingredient(s))
- Drug product patents, which include formulation/composition patents
- Method-of-use patents that claim one or more approved methods of using the approved drug product

This information, as provided by the sponsor on Form FDA 3542, will be published 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.

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 deletions, the [Orange Book](#), updated daily, should be consulted for the most recent patent and exclusivity information.

¹ 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).

² See 21 CFR 314.50(i) (4) and 314.94(a) (12) (vi). The submission date for patent information is determined in accordance with 21 CFR 314.53(d) (5).

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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			
	9242986	Dec 08, 2029	DS DP			
<u>ABALOPARATIDE - TYMLOS</u>						
N 208743 001	7803770	Mar 26, 2028		U-2009	NCE	Apr 28, 2022
	8148333	Nov 08, 2027	DP			
	8748382	Oct 03, 2027		U-2009		
<u>ABEMACICLIB - VERZENIO</u>						
N 208716 001	7855211	Dec 15, 2029	DS DP U-2132		I-768	Feb 26, 2021
	7855211	Dec 15, 2029	DS DP U-2135		NCE	Sep 28, 2022
	7855211	Dec 15, 2029	DS DP U-2251			
<u>ABEMACICLIB - VERZENIO</u>						
N 208716 002	7855211	Dec 15, 2029	DS DP U-2132		I-768	Feb 26, 2021
	7855211	Dec 15, 2029	DS DP U-2135		NCE	Sep 28, 2022
	7855211	Dec 15, 2029	DS DP U-2251			
<u>ABEMACICLIB - VERZENIO</u>						
N 208716 003	7855211	Dec 15, 2029	DS DP U-2132		I-768	Feb 26, 2021
	7855211	Dec 15, 2029	DS DP U-2135		NCE	Sep 28, 2022
	7855211	Dec 15, 2029	DS DP U-2251			
<u>ABEMACICLIB - VERZENIO</u>						
N 208716 004	7855211	Dec 15, 2029	DS DP U-1981		I-768	Feb 26, 2021
	7855211	Dec 15, 2029	DS DP U-2132		NCE	Sep 28, 2022
	7855211	Dec 15, 2029	DS DP U-2135			
	7855211	Dec 15, 2029	DS DP U-2251			
<u>ABIRATERONE ACETATE - ZYTIGA</u>						
N 202379 001					I-765	Feb 07, 2021
<u>ABIRATERONE ACETATE - ZYTIGA</u>						
N 202379 002	8822438	Aug 24, 2027		U-1579	Y	Feb 07, 2021
	8822438	Aug 24, 2027		U-1580	Y	
	8822438	Aug 24, 2027		U-2235	Y	
<u>ABIRATERONE ACETATE - YONSA</u>						
N 210308 001	10292990	May 20, 2034		U-2535		
	9889144	Mar 17, 2034	DP			
<u>ACALABRUTINIB - CALOQUENCE</u>						
N 210259 001	10167291	Jul 01, 2036		DP U-2145	I-817	Nov 21, 2022
	10167291	Jul 01, 2036		DP U-2666	NCE	Oct 31, 2022
	10167291	Jul 01, 2036		DP U-2667	ODE-175	Oct 31, 2024
	10167291	Jul 01, 2036		DP U-2668	ODE-274	Nov 21, 2026
	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		
	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		
	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		

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	10383834	Nov 13, 2028	U-2262		M-196	Jan 27, 2020
	10383834	Nov 13, 2028	U-2621		PED	Jul 27, 2020
	6992218	Jun 06, 2021	DP			
	6992218*PED	Dec 06, 2021				
	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			
	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; BUTALBITAL - BUTALBITAL AND ACETAMINOPHEN</u>						
A 213115 001					CGT	Jul 28, 2020
<u>ACETAMINOPHEN; OXYCODONE HYDROCHLORIDE - XARTEMIS XR</u>						
N 204031 001	6488962	Jun 20, 2020	DP			
	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			
	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			

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APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<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>ACLIDINIUM BROMIDE - TUDORZA PRESSAIR</u>						
N 202450	001	10034867	Jul 07, 2020	DP U-2513		
		10034867	Jul 07, 2020	DP U-2514		
		10085974	Mar 13, 2029	DP U-2513		
		6681768	Aug 07, 2022	DP		
		7078412	Jul 07, 2020	DS DP U-2513		
		8051851	Apr 22, 2027	DP		
		9056100	Jul 07, 2020	DP U-2513		
		9333195	Jul 07, 2020	DP U-2513		
		RE46417	Feb 10, 2025	DS DP U-2513		
<u>ACLIDINIUM BROMIDE; FORMOTEROL FUMARATE - DUAKLIR PRESSAIR</u>						
N 210595	001	10034867	Jul 07, 2020	DP U-2513	NC	Mar 29, 2022
		10034867	Jul 07, 2020	DP U-2514		
		10085974	Mar 13, 2029	DP U-2513		
		6681768	Aug 07, 2022	DP		
		7078412	Jul 07, 2020	DS DP U-2513		
		7750023	Jul 07, 2020	DP U-2513		
		8051851	Apr 22, 2027	DP		
		8129405	Jul 07, 2020	DP U-2513		
		9056100	Jul 07, 2020	DP U-2513		
		9333195	Jul 07, 2020	DP U-2513		
		RE46417	Feb 10, 2025	DS DP U-2513		
<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>ACYCLOVIR; HYDROCORTISONE - XERESE</u>						
N 022436	001	7223387	Nov 13, 2022	DP U-1006		
		7223387	Nov 13, 2022	DP U-1484		
<u>ADAPALENE - DIFFERIN</u>						
N 021753	001	7579377	Feb 23, 2025	U-818		
		7737181	Aug 29, 2024	DP		
		7834060	Mar 12, 2023	U-1078		
		7838558	Mar 12, 2023	DP		
		7868044	Mar 12, 2023	U-1078		
		8703820	Mar 12, 2023	U-1078		
<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		
		7964202	Sep 01, 2024	DP U-1078		
		8071644	Jul 18, 2027	DP U-1078		
		8080537	Jul 18, 2027	U-1078		
		8105618	Dec 23, 2022	U-1078		
		8129362	Jul 18, 2027	U-1078		
		8241649	Dec 23, 2022	DP		
		8445543	Jul 12, 2027	U-1078		
		8809305	Dec 23, 2022	U-1078		
		8936800	Dec 23, 2022	DP U-1078		
<u>ADAPALENE; BENZOYL PEROXIDE - EPIDUO FORTE</u>						
N 207917	001	8445543	Dec 23, 2022	U-1078		
		8703820	Mar 12, 2023	U-1078		
		8729127	Mar 12, 2023	U-1078		
		8785420	Dec 23, 2022	U-1078		

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<u>ADAPALENE; BENZOYL PEROXIDE - EPIDUO FORTE</u>						
N 207917	001	8809305				
		8936800				
		9381179				
		9387187				
		9814690				
<u>AFAMELANOTIDE - SCENESSE</u>						
N 210797	001	10076555				
		8334265				
<u>AFATINIB DIMALEATE - GILOTRIF</u>						
N 201292	001	10004743				
		8426586				
		8545884				
		9539258				
		RE43431				
<u>AFATINIB DIMALEATE - GILOTRIF</u>						
N 201292	002	10004743				
		8426586				
		8545884				
		9539258				
		RE43431				
<u>AFATINIB DIMALEATE - GILOTRIF</u>						
N 201292	003	10004743				
		8426586				
		8545884				
		9539258				
		RE43431				
<u>AIR POLYMER-TYPE A - EXEM FOAM KIT</u>						
N 212279	001	9034300				
		9259494				
		9849199				
<u>ALBUMIN HUMAN - OPTISON</u>						
N 020899	001	6723303				
<u>ALBUTEROL SULFATE - ACCUNEB</u>						
N 020949	001	6702997				
<u>ALBUTEROL SULFATE - ACCUNEB</u>						
N 020949	002	6702997				
<u>ALBUTEROL SULFATE - VENTOLIN HFA</u>						
N 020983	001	7500444				
		7500444*PED				
		7832351				
		9861771				
<u>ALBUTEROL SULFATE - PROAIR HFA</u>						
N 021457	001	10022509				
		10022510				
		10086156				
		7105152				
		8132712				
		9463289				
		9808587				
<u>ALBUTEROL SULFATE - PROAIR RESPICLICK</u>						
N 205636	001	10022510				
		10124131				
		6701917				
		6718972				
		6748947				
		6871646				
		7540282				
		8006690				
		8651103				

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<u>ALBUTEROL SULFATE - PROAIR RESPICLICK</u>						
N 205636 001	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			
	6701917	Jun 23, 2021	DP			
	6718972	Jun 23, 2021	DP			
	6748947	Jun 23, 2021	DP			
	6871646	Jun 23, 2021	DP			
	7540282	May 06, 2023	DP			
	8006690	Jun 23, 2021	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; IPRATROPIUM BROMIDE - DUONEB</u>						
N 020950 001	6632842	Dec 28, 2021		U-532		
<u>ALBUTEROL SULFATE; IPRATROPIUM BROMIDE - COMBIVENT RESPIMAT</u>						
N 021747 001	6988496	Feb 23, 2020	DP			
	7284474	Aug 26, 2024	DP			
	7396341	Oct 10, 2026	DP			
	7837235	Mar 13, 2028	DP			
	7896264	May 26, 2025	DP			
	7988001	Aug 04, 2021	DP			
	8733341	Oct 16, 2030	DP			
	9027967	Mar 31, 2027	DP			
<u>ALCAFTADINE - LASTACFT</u>						
N 022134 001	8664215	Dec 23, 2027		U-1493		
<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	9126931	May 29, 2031	DS		I-756	Nov 06, 2020
	9365514	Mar 04, 2032	DP		NCE	Dec 11, 2020
	9440922	Jun 09, 2030	DP		ODE-105	Dec 11, 2022
					ODE-159	Nov 06, 2024
<u>ALENDRONATE SODIUM - BINOSTO</u>						
N 202344 001	7488496	Aug 11, 2023	DS DP			
	7964212	Mar 06, 2023	DS DP			
<u>ALISKIREN HEMIFUMARATE - TEKTRUNA</u>						
N 021985 001	8617595	Feb 19, 2026	DP			
	8617595*PED	Aug 19, 2026				
<u>ALISKIREN HEMIFUMARATE - TEKTRUNA</u>						
N 021985 002	8617595	Feb 19, 2026	DP			
	8617595*PED	Aug 19, 2026				
<u>ALISKIREN HEMIFUMARATE - TEKTRUNA</u>						
N 210709 001					NP PED	Nov 14, 2020 May 14, 2021
<u>ALISKIREN HEMIFUMARATE; AMLODIPINE BESYLATE - TEKAMLO</u>						
N 022545 001	8613949	Dec 21, 2029	DP			

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<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; AMLODIPINE BESYLATE; HYDROCHLOROTHIAZIDE - AMTURNIDE</u>						
N 200045	001	8183295	May 16, 2023	DP		
		8618174	Nov 15, 2021	DP		
<u>ALISKIREN HEMIFUMARATE; AMLODIPINE BESYLATE; HYDROCHLOROTHIAZIDE - AMTURNIDE</u>						
N 200045	002	8183295	May 16, 2023	DP		
		8618174	Nov 15, 2021	DP		
<u>ALISKIREN HEMIFUMARATE; AMLODIPINE BESYLATE; HYDROCHLOROTHIAZIDE - AMTURNIDE</u>						
N 200045	003	8183295	May 16, 2023	DP		
		8618174	Nov 15, 2021	DP		
<u>ALISKIREN HEMIFUMARATE; AMLODIPINE BESYLATE; HYDROCHLOROTHIAZIDE - AMTURNIDE</u>						
N 200045	004	8183295	May 16, 2023	DP		
		8618174	Nov 15, 2021	DP		
<u>ALISKIREN HEMIFUMARATE; AMLODIPINE BESYLATE; HYDROCHLOROTHIAZIDE - AMTURNIDE</u>						
N 200045	005	8183295	May 16, 2023	DP		
		8618174	Nov 15, 2021	DP		
<u>ALISKIREN HEMIFUMARATE; HYDROCHLOROTHIAZIDE - TEKTURNA HCT</u>						
N 022107	001	8618172	Jul 13, 2028	DP		
		9023893	Mar 03, 2022	DP		
<u>ALISKIREN HEMIFUMARATE; HYDROCHLOROTHIAZIDE - TEKTURNA HCT</u>						
N 022107	002	8618172	Jul 13, 2028	DP		
		9023893	Mar 03, 2022	DP		
<u>ALISKIREN HEMIFUMARATE; HYDROCHLOROTHIAZIDE - TEKTURNA HCT</u>						
N 022107	003	8618172	Jul 13, 2028	DP		
		9023893	Mar 03, 2022	DP		
<u>ALISKIREN HEMIFUMARATE; HYDROCHLOROTHIAZIDE - TEKTURNA HCT</u>						
N 022107	004	8618172	Jul 13, 2028	DP		
		9023893	Mar 03, 2022	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	Nov 26, 2028		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	Nov 26, 2028		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

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<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	Nov 26, 2028		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	
<u>ALOGLIPTIN BENZOATE - NESINA</u>						
N 022271	002	7807689	Jun 27, 2028	DS DP		U-1337
		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
		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
		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
		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	6329404	Jun 19, 2021		DP	U-1334
		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	6329404	Jun 19, 2021		DP	U-1334
		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	6329404	Jun 19, 2021		DP	U-1334
		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	6329404	Jun 19, 2021		DP	U-1334
		7807689	Jun 27, 2028	DS DP		U-1337
		8173663	Mar 15, 2025			U-1338
		8288539	Mar 15, 2025	DS		
		8637079	Jun 04, 2029		DP	

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<u>ALOGLIPTIN BENZOATE; PIOGLITAZONE HYDROCHLORIDE - OSENI</u>						
N 022426 005	6329404	Jun 19, 2021	DP U-1334			
	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 006	6329404	Jun 19, 2021	DP U-1334			
	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	Sep 28, 2030	DS DP U-2539		NCE	May 24, 2024
	8476268	Sep 10, 2029	DS DP			
<u>ALPELISIB - PIORAY</u>						
N 212526 002	8227462	Sep 28, 2030	DS DP U-2539		NCE	May 24, 2024
	8476268	Sep 10, 2029	DS DP			
<u>ALPELISIB - PIORAY</u>						
N 212526 003	8227462	Sep 28, 2030	DS DP U-2539		NCE	May 24, 2024
	8476268	Sep 10, 2029	DS DP			
<u>ALVIMOPAN - ENTEREG</u>						
N 021775 001	6469030	Nov 29, 2020	U-879			
	8112290	Jul 31, 2030	U-1443	Y		
	8645160	Jun 18, 2029	U-1485	Y		
	8946262	Feb 12, 2030	U-1655			
<u>AMANTADINE HYDROCHLORIDE - GOCOVRI</u>						
N 208944 001	10154971	Dec 04, 2034	U-2459		I-769	Aug 24, 2020
	8389578	Jan 22, 2028	U-2105		ODE-153	Aug 24, 2024
	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 - GOCOVRI</u>						
N 208944 002	10154971	Dec 04, 2034	U-2459		I-769	Aug 24, 2020
	8389578	Jan 22, 2028	U-2105		ODE-153	Aug 24, 2024
	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			

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<u>AMANTADINE HYDROCHLORIDE - OSMOLEX ER</u>						
N 209410	001	8252331	Mar 13, 2030	DP		
		8574626	Nov 28, 2025	DP U-20		
<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	
		8252331	Mar 13, 2030	DP		
		8574626	Nov 28, 2025	DP U-20		
<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		
		8574626	Nov 28, 2025	DP U-20		
<u>AMBRISENTAN - LETAIRIS</u>						
N 022081	001	8377933	Dec 11, 2027		U-1754	
		9474752	Dec 11, 2027		U-1754	
		9549926	Oct 14, 2031		U-1965	
<u>AMBRISENTAN - 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 - RUZURGI</u>						
N 209321	001				ODE-244	May 06, 2026
<u>AMIFAMPRIDINE PHOSPHATE - FIRDAPSE</u>						
N 208078	001				NCE	Nov 28, 2023
					ODE-223	Nov 28, 2025
<u>AMIKACIN SULFATE - ARIKAYCE KIT</u>						
N 207356	001	10251900	May 15, 2035		U-2414	
		7718189	Jun 06, 2025	DP	U-2415	Sep 28, 2025
		8226975	Aug 15, 2028	DP		Sep 28, 2030
		8632804	Dec 05, 2026		U-2416	
		8642075	Dec 05, 2026	DP		
		8679532	Dec 05, 2026		U-2415	
		8802137	Apr 08, 2024	DP	U-2414	
		9566234	Jan 18, 2034	DP	U-2415	
		9827317	Apr 08, 2024	DP	U-2415	
		9895385	May 15, 2035		U-2417	
<u>AMINOCAPROIC ACID - AMINOCAPROIC ACID</u>						
A 212492	001				CGT	May 30, 2020
<u>AMINOCAPROIC ACID - AMINOCAPROIC ACID</u>						
A 212780	001				CGT	Feb 26, 2020
<u>AMINOLEVULINIC ACID HYDROCHLORIDE - LEVULAN</u>						
N 020965	001	10357567	Jan 12, 2038		U-804	Mar 09, 2021
<u>AMINOLEVULINIC ACID HYDROCHLORIDE - GLEOLAN</u>						
N 208630	001				ODE-146	Jun 06, 2024
<u>AMIODARONE HYDROCHLORIDE - NEXTERONE</u>						
N 022325	001	6869939	May 04, 2022	DP		
		7635773	Mar 13, 2029	DP		

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<u>AMIODARONE HYDROCHLORIDE - NEXTERONE</u>						
N 022325	002	6869939	May 04, 2022	DP		
		7635773	Mar 13, 2029	DP		
<u>AMIODARONE HYDROCHLORIDE - NEXTERONE</u>						
N 022325	003	6869939	May 04, 2022	DP		
		7635773	Mar 13, 2029	DP		
<u>AMISULPRIDE - BARHEMSYS</u>						
N 209510	001				NCE	Feb 26, 2025
<u>AMLODIPINE BESYLATE - AMLODIPINE BESYLATE</u>						
N 022026	001	6828339	Nov 20, 2022	DS		
<u>AMLODIPINE BESYLATE - AMLODIPINE BESYLATE</u>						
N 022026	002	6828339	Nov 20, 2022	DS		
<u>AMLODIPINE BESYLATE - AMLODIPINE BESYLATE</u>						
N 022026	003	6828339	Nov 20, 2022	DS		
<u>AMLODIPINE BESYLATE; CELECOXIB - CONSENSI</u>						
N 210045	001	10350171	Jun 14, 2038	DP	NC	May 31, 2021
		9662315	May 22, 2029	DP U-2410		
<u>AMLODIPINE BESYLATE; CELECOXIB - CONSENSI</u>						
N 210045	002	10350171	Jun 14, 2038	DP	NC	May 31, 2021
		9662315	May 22, 2029	DP U-2410		
<u>AMLODIPINE BESYLATE; CELECOXIB - CONSENSI</u>						
N 210045	003	10350171	Jun 14, 2038	DP	NC	May 31, 2021
		9662315	May 22, 2029	DP U-2410		
<u>AMLODIPINE BESYLATE; HYDROCHLOROTHIAZIDE; VALSARTAN - EXFORGE HCT</u>						
N 022314	001	8101599	May 16, 2023	DP		
		8475839	May 16, 2023	DP		
		8475839*PED	Nov 16, 2023			
<u>AMLODIPINE BESYLATE; HYDROCHLOROTHIAZIDE; VALSARTAN - EXFORGE HCT</u>						
N 022314	002	8101599	May 16, 2023	DP		
		8475839	May 16, 2023	DP		
		8475839*PED	Nov 16, 2023			
<u>AMLODIPINE BESYLATE; HYDROCHLOROTHIAZIDE; VALSARTAN - EXFORGE HCT</u>						
N 022314	003	8101599	May 16, 2023	DP		
		8475839	May 16, 2023	DP		
		8475839*PED	Nov 16, 2023			
<u>AMLODIPINE BESYLATE; HYDROCHLOROTHIAZIDE; VALSARTAN - EXFORGE HCT</u>						
N 022314	004	8101599	May 16, 2023	DP		
		8475839	May 16, 2023	DP		
		8475839*PED	Nov 16, 2023			
<u>AMLODIPINE BESYLATE; HYDROCHLOROTHIAZIDE; VALSARTAN - EXFORGE HCT</u>						
N 022314	005	8101599	May 16, 2023	DP		
		8475839	May 16, 2023	DP		
		8475839*PED	Nov 16, 2023			
<u>AMLODIPINE BESYLATE; PERINDOPRIL ARGININE - PRESTALIA</u>						
N 205003	001	6696481	Apr 15, 2023	DS DP U-3		
		7846961	Oct 05, 2029	DS DP U-3		
<u>AMLODIPINE BESYLATE; PERINDOPRIL ARGININE - PRESTALIA</u>						
N 205003	002	6696481	Apr 15, 2023	DS DP U-3		
		7846961	Oct 05, 2029	DS DP U-3		
<u>AMLODIPINE BESYLATE; PERINDOPRIL ARGININE - PRESTALIA</u>						
N 205003	003	6696481	Apr 15, 2023	DS DP U-3		
		7846961	Oct 05, 2029	DS DP U-3		
<u>AMOXICILLIN - MOXATAG</u>						
N 050813	001	6544555	Oct 13, 2020	DS DP U-897		
		6669948	Oct 13, 2020	DS DP U-897		

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<u>AMOXICILLIN - MOXATAG</u>						
N 050813	001	6723341	Oct 13, 2020	DS DP	U-897	
		8299052	May 07, 2027		U-1304	
		8357394	Dec 08, 2026	DP		
		8778924	Dec 08, 2026	DS DP	U-897	
<u>AMOXICILLIN; CLAVULANATE POTASSIUM - AUGMENTIN XR</u>						
N 050785	001	6746692	Apr 04, 2020	DP		
		6783773	Apr 04, 2020	DP		
		6878386	Apr 04, 2020		U-926	
		7217430	Apr 04, 2020	DP	U-926	
		7250176	Apr 04, 2020		U-926	
<u>AMOXICILLIN; OMEPRAZOLE MAGNESIUM; RIFABUTIN - TALICIA</u>						
N 213004	001	10238606	Feb 12, 2034	DP		
		9050263	Feb 12, 2034	DP	U-2660	
		9498445	Feb 12, 2034	DP	U-2660	
		9603806	Feb 12, 2034	DP	U-2660	
<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		
<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		
<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		
<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		
<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		
<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		
<u>AMPHETAMINE - DYANA VEL XR</u>						
N 208147	001	10086087	Mar 15, 2027	DP		
		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 ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATE - ADDERALL 10</u>						
N 011522	007	6384020	Jul 06, 2020			

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<u>AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATE - ADDERALL 20</u>						
N 011522	008	6384020	Jul 06, 2020			
<u>AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATE - ADDERALL 5</u>						
N 011522	009	6384020	Jul 06, 2020			
<u>AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATE - ADDERALL 30</u>						
N 011522	010	6384020	Jul 06, 2020			
<u>AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATE - ADDERALL 7.5</u>						
N 011522	011	6384020	Jul 06, 2020			
<u>AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATE - ADDERALL 12.5</u>						
N 011522	012	6384020	Jul 06, 2020			
<u>AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATE - ADDERALL 15</u>						
N 011522	013	6384020	Jul 06, 2020			
<u>AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATE - MYDAYIS</u>						
N 022063	001	6913768	May 24, 2023	DP U-2025	M-248	Sep 13, 2022
		8846100	Aug 24, 2029	DP	NP	Jun 20, 2020
		9173857	May 12, 2026	U-2025	PED	Mar 13, 2023
<u>AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATE - MYDAYIS</u>						
N 022063	002	6913768	May 24, 2023	DP U-2025	M-248	Sep 13, 2022
		8846100	Aug 24, 2029	DP	NP	Jun 20, 2020
		9173857	May 12, 2026	U-2025	PED	Mar 13, 2023
<u>AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATE - MYDAYIS</u>						
N 022063	003	6913768	May 24, 2023	DP U-2025	M-248	Sep 13, 2022
		8846100	Aug 24, 2029	DP	NP	Jun 20, 2020
		9173857	May 12, 2026	U-2025	PED	Mar 13, 2023
<u>AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATE - MYDAYIS</u>						
N 022063	004	6913768	May 24, 2023	DP U-2025	M-248	Sep 13, 2022
		8846100	Aug 24, 2029	DP	NP	Jun 20, 2020
		9173857	May 12, 2026	U-2025	PED	Mar 13, 2023
<u>AMPHETAMINE SULFATE - EVEKEO ODT</u>						
N 209905	001	10130580	Apr 19, 2024	DP		
		10441554	Mar 10, 2037	DP		
<u>AMPHETAMINE SULFATE - EVEKEO ODT</u>						
N 209905	002	10130580	Apr 19, 2024	DP		
		10441554	Mar 10, 2037	DP		
<u>AMPHETAMINE SULFATE - EVEKEO ODT</u>						
N 209905	003	10130580	Apr 19, 2024	DP		
		10441554	Mar 10, 2037	DP		
<u>AMPHETAMINE SULFATE - EVEKEO ODT</u>						
N 209905	004	10130580	Apr 19, 2024	DP		
		10441554	Mar 10, 2037	DP		
<u>ANGIOTENSIN II ACETATE - GIAPREZA</u>						
N 209360	001	10028995	Dec 18, 2034	U-2338	NCE	Dec 21, 2022
		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		
		9220745	Dec 18, 2034	U-2217		
		9220745	Dec 18, 2034	U-2218		

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<u>ANGIOTENSIN II ACETATE - GIAPREZA</u>						
N 209360	001	9572856	Nov 20, 2030	U-2221		
		9867863	Dec 16, 2029	U-2231		
<u>ANGIOTENSIN II ACETATE - GIAPREZA</u>						
N 209360	002	10028995	Dec 18, 2034	U-2338	NCE	Dec 21, 2022
		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		
		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>ANIDULAFUNGIN - ERAXIS</u>						
N 021632	001	5965525	Feb 17, 2020	DS DP U-540		
		6960564	Apr 12, 2021	DP U-540		
		7709444	Apr 12, 2021	DP U-540		
<u>ANIDULAFUNGIN - ERAXIS</u>						
N 021632	002	5965525	Feb 17, 2020	DS DP U-540		
		6960564	Apr 12, 2021	DP U-540		
		7709444	Apr 12, 2021	DP U-540		
<u>APALUTAMIDE - ERLEADA</u>						
N 210951	001	10052314	Sep 23, 2033	U-2381	I-808	Sep 17, 2022
		10052314	Sep 23, 2033	U-2382	NCE	Feb 14, 2023
		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		
<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>APREMILAST - OTEZLA</u>						
N 205437	001	10092541	May 29, 2034	U-2403	I-803	Jul 19, 2022
		10092541	May 29, 2034	U-2659	ODE-248	Jul 19, 2026
		6962940	Mar 19, 2023	U-1504		
		6962940	Mar 19, 2023	U-2656		
		6962940	Mar 19, 2023	U-2657		
		6962940	Mar 19, 2023	U-2658		
		7208516	Mar 19, 2023	U-1505		
		7427638	Feb 16, 2028	DS DP		
		7659302	Mar 19, 2023	U-1505		
		7659302	Mar 19, 2023	U-1595		
		7659302	Mar 19, 2023	U-2657		

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<u>APREMILAST - OTEZLA</u>						
N 205437 001	7659302	Mar 19, 2023				
	7893101	Dec 09, 2023	DS DP			
	8455536	Mar 19, 2023			U-1505	
	8455536	Mar 19, 2023			U-1595	
	8455536	Mar 19, 2023			U-2657	
	8455536	Mar 19, 2023			U-2658	
	8802717	Mar 19, 2023			U-1561	
	9018243	Mar 19, 2023			U-1505	
	9018243	Mar 19, 2023			U-1595	
	9018243	Mar 19, 2023			U-2656	
	9018243	Mar 19, 2023			U-2657	
	9018243	Mar 19, 2023			U-2658	
	9724330	Mar 19, 2023			U-1561	
	9724330	Mar 19, 2023			U-1595	
	9724330	Mar 19, 2023			U-2656	
	9724330	Mar 19, 2023			U-2657	
	9724330	Mar 19, 2023			U-2658	
	9872854	May 29, 2034			U-2232	
	9872854	May 29, 2034			U-2233	
<u>APREMILAST - OTEZLA</u>						
N 205437 002	10092541	May 29, 2034			I-803	Jul 19, 2022
	10092541	May 29, 2034			ODE-248	Jul 19, 2026
	6962940	Mar 19, 2023			U-1504	
	6962940	Mar 19, 2023			U-2656	
	6962940	Mar 19, 2023			U-2657	
	6962940	Mar 19, 2023			U-2658	
	7208516	Mar 19, 2023			U-1505	
	7427638	Feb 16, 2028	DS DP			
	7659302	Mar 19, 2023			U-1505	
	7659302	Mar 19, 2023			U-1595	
	7659302	Mar 19, 2023			U-2657	
	7659302	Mar 19, 2023			U-2658	
	7893101	Dec 09, 2023	DS DP			
	8455536	Mar 19, 2023			U-1505	
	8455536	Mar 19, 2023			U-1595	
	8455536	Mar 19, 2023			U-2657	
	8455536	Mar 19, 2023			U-2658	
	8802717	Mar 19, 2023			U-1561	
	9018243	Mar 19, 2023			U-1505	
	9018243	Mar 19, 2023			U-1595	
	9018243	Mar 19, 2023			U-2656	
	9018243	Mar 19, 2023			U-2657	
	9018243	Mar 19, 2023			U-2658	
	9724330	Mar 19, 2023			U-1561	
	9724330	Mar 19, 2023			U-1595	
	9724330	Mar 19, 2023			U-2656	
	9724330	Mar 19, 2023			U-2657	
	9724330	Mar 19, 2023			U-2658	
	9872854	May 29, 2034			U-2232	
	9872854	May 29, 2034			U-2233	
<u>APREMILAST - OTEZLA</u>						
N 205437 003	10092541	May 29, 2034			I-803	Jul 19, 2022
	10092541	May 29, 2034			ODE-248	Jul 19, 2026
	6962940	Mar 19, 2023			U-1504	
	6962940	Mar 19, 2023			U-2656	
	6962940	Mar 19, 2023			U-2657	
	6962940	Mar 19, 2023			U-2658	
	7208516	Mar 19, 2023			U-1505	
	7427638	Feb 16, 2028	DS DP			
	7659302	Mar 19, 2023			U-1505	
	7659302	Mar 19, 2023			U-1595	
	7659302	Mar 19, 2023			U-2657	
	7659302	Mar 19, 2023			U-2658	
	7893101	Dec 09, 2023	DS DP			
	8455536	Mar 19, 2023			U-1505	
	8455536	Mar 19, 2023			U-1595	
	8455536	Mar 19, 2023			U-2657	

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<u>APREMILAST - OTEZLA</u>						
N 205437	003 8455536	Mar 19, 2023	U-2658			
	8802717	Mar 19, 2023	U-1561			
	9018243	Mar 19, 2023	U-1505			
	9018243	Mar 19, 2023	U-1595			
	9018243	Mar 19, 2023	U-2656			
	9018243	Mar 19, 2023	U-2657			
	9018243	Mar 19, 2023	U-2658			
	9724330	Mar 19, 2023	U-1561			
	9724330	Mar 19, 2023	U-1595			
	9724330	Mar 19, 2023	U-2656			
	9724330	Mar 19, 2023	U-2657			
	9724330	Mar 19, 2023	U-2658			
	9872854	May 29, 2034	U-2232			
	9872854	May 29, 2034	U-2233			
<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			
<u>APREPITANT - CINVANTI</u>						
N 209296	001 10500208	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>ARFORMOTEROL TARTRATE - BROVANA</u>						
N 021912	001 6472563	Nov 09, 2021	DS			
	6667344	Jun 22, 2021	DP			
	6720453	Nov 09, 2021	DS			
	6814953	Jun 22, 2021	U-793			
	7145036	Nov 09, 2021	DS			
	7348362	Jun 22, 2021	DP U-793			
	7462645	Jun 22, 2021	U-793			
	7465756	Jun 22, 2021	DP			
	7473710	Jun 22, 2021	U-793			
	7541385	Jun 22, 2021	U-793			
	8110706	Nov 09, 2021	DP			
<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>ARIPIPRAZOLE - ABILIFY</u>						
N 021436	001 7053092	Jan 28, 2022	U-839		ODE-80	Dec 12, 2021
	8017615	Jun 16, 2024	DP			
	8017615*PED	Dec 16, 2024				
	8580796	Sep 25, 2022	DS			
	8580796*PED	Mar 25, 2023				
	8642600	Jan 28, 2022	U-1492			
	8642600*PED	Jul 28, 2022				
	8642760	Sep 25, 2022	DS			
	8642760*PED	Mar 25, 2023				
	8759350	Mar 02, 2027	U-1529			
	9089567	Jan 28, 2022	U-543			
	9125939	Jul 28, 2026	U-1749			
	9359302	Sep 25, 2022	DS DP U-1859			
	9387182	Dec 25, 2023	U-1529			

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<u>ARIPIRAZOLE - ABILIFY</u>						
N 021436 001	7053092	Jan 28, 2022		U-839	ODE-80	Dec 12, 2021
	8017615	Jun 16, 2024		DP		
	8017615*PED	Dec 16, 2024				
	8580796	Sep 25, 2022	DS			
	8580796*PED	Mar 25, 2023				
	8642600	Jan 28, 2022		U-1492		
	8642600*PED	Jul 28, 2022				
	8642760	Sep 25, 2022	DS			
	8642760*PED	Mar 25, 2023				
	8759350	Mar 02, 2027		U-1529		
	9089567	Jan 28, 2022		U-543		
	9125939	Jul 28, 2026		U-1749		
	9359302	Sep 25, 2022	DS DP	U-1859		
	9387182	Dec 25, 2023		U-1529		
<u>ARIPIRAZOLE - ABILIFY</u>						
N 021436 002	7053092	Jan 28, 2022		U-839	ODE-80	Dec 12, 2021
	8017615	Jun 16, 2024		DP		
	8017615*PED	Dec 16, 2024				
	8580796	Sep 25, 2022	DS			
	8580796*PED	Mar 25, 2023				
	8642600	Jan 28, 2022		U-1492		
	8642600*PED	Jul 28, 2022				
	8642760	Sep 25, 2022	DS			
	8642760*PED	Mar 25, 2023				
	8759350	Mar 02, 2027		U-1529		
	9089567	Jan 28, 2022		U-543		
	9125939	Jul 28, 2026		U-1749		
	9359302	Sep 25, 2022	DS DP	U-1859		
	9387182	Dec 25, 2023		U-1529		
<u>ARIPIRAZOLE - ABILIFY</u>						
N 021436 003	7053092	Jan 28, 2022		U-839	ODE-80	Dec 12, 2021
	8017615	Jun 16, 2024		DP		
	8017615*PED	Dec 16, 2024				
	8580796	Sep 25, 2022	DS			
	8580796*PED	Mar 25, 2023				
	8642600	Jan 28, 2022		U-1492		
	8642600*PED	Jul 28, 2022				
	8642760	Sep 25, 2022	DS			
	8642760*PED	Mar 25, 2023				
	8759350	Mar 02, 2027		U-1529		
	9089567	Jan 28, 2022		U-543		
	9125939	Jul 28, 2026		U-1749		
	9359302	Sep 25, 2022	DS DP	U-1859		
	9387182	Dec 25, 2023		U-1529		
<u>ARIPIRAZOLE - ABILIFY</u>						
N 021436 004	7053092	Jan 28, 2022		U-839	ODE-80	Dec 12, 2021
	8017615	Jun 16, 2024		DP		
	8017615*PED	Dec 16, 2024				
	8580796	Sep 25, 2022	DS			
	8580796*PED	Mar 25, 2023				
	8642600	Jan 28, 2022		U-1492		
	8642600*PED	Jul 28, 2022				
	8642760	Sep 25, 2022	DS			
	8642760*PED	Mar 25, 2023				
	8759350	Mar 02, 2027		U-1529		
	9089567	Jan 28, 2022		U-543		
	9125939	Jul 28, 2026		U-1749		
	9359302	Sep 25, 2022	DS DP	U-1859		
	9387182	Dec 25, 2023		U-1529		
<u>ARIPIRAZOLE - ABILIFY</u>						
N 021436 005	7053092	Jan 28, 2022		U-839	ODE-80	Dec 12, 2021
	8017615	Jun 16, 2024		DP		
	8017615*PED	Dec 16, 2024				
	8580796	Sep 25, 2022	DS			
	8580796*PED	Mar 25, 2023				
	8642600	Jan 28, 2022		U-1492		

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APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>ARIPIRAZOLE - ABILIFY</u>						
N 021436 005	8642600*PED	Jul 28, 2022				
	8642760	Sep 25, 2022	DS			
	8642760*PED	Mar 25, 2023				
	8759350	Mar 02, 2027		U-1529		
	9089567	Jan 28, 2022		U-543		
	9125939	Jul 28, 2026		U-1749		
	9359302	Sep 25, 2022	DS DP	U-1859		
	9387182	Dec 25, 2023		U-1529		
<u>ARIPIRAZOLE - ABILIFY</u>						
N 021436 006	7053092	Jan 28, 2022		U-839	ODE-80	Dec 12, 2021
	8017615	Jun 16, 2024	DP			
	8017615*PED	Dec 16, 2024				
	8580796	Sep 25, 2022	DS			
	8580796*PED	Mar 25, 2023				
	8642600	Jan 28, 2022		U-1492		
	8642600*PED	Jul 28, 2022				
	8642760	Sep 25, 2022	DS			
	8642760*PED	Mar 25, 2023				
	8759350	Mar 02, 2027		U-1529		
	9089567	Jan 28, 2022		U-543		
	9125939	Jul 28, 2026		U-1749		
	9359302	Sep 25, 2022	DS DP	U-1859		
	9387182	Dec 25, 2023		U-1529		
<u>ARIPIRAZOLE - ABILIFY</u>						
N 021713 001	6977257	Apr 24, 2022	DP		ODE-80	Dec 12, 2021
	6977257*PED	Oct 24, 2022				
	7053092	Jan 28, 2022		U-839		
	8642600	Jan 28, 2022		U-1492		
	8642600*PED	Jul 28, 2022				
	8759350	Mar 02, 2027		U-1529		
	9387182	Dec 25, 2023		U-1529		
<u>ARIPIRAZOLE - ABILIFY</u>						
N 021729 002	7053092	Jan 28, 2022		U-839	ODE-80	Dec 12, 2021
	8017615	Jun 16, 2024	DP			
	8017615*PED	Dec 16, 2024				
	8518421	Jan 24, 2021	DP			
	8518421*PED	Jul 24, 2021				
	8580796	Sep 25, 2022	DS			
	8580796*PED	Mar 25, 2023				
	8642600	Jan 28, 2022		U-1492		
	8642600*PED	Jul 28, 2022				
	8642760	Sep 25, 2022	DS			
	8642760*PED	Mar 25, 2023				
	8759350	Mar 02, 2027		U-1529		
	9089567	Jan 28, 2022		U-543		
	9125939	Jul 28, 2026		U-1749		
	9358207	Apr 12, 2020	DP			
	9359302	Sep 25, 2022	DS DP	U-1859		
	9387182	Dec 25, 2023		U-1529		
<u>ARIPIRAZOLE - ABILIFY</u>						
N 021729 003	7053092	Jan 28, 2022		U-839	ODE-80	Dec 12, 2021
	8017615	Jun 16, 2024	DP			
	8017615*PED	Dec 16, 2024				
	8518421	Jan 24, 2021	DP			
	8518421*PED	Jul 24, 2021				
	8580796	Sep 25, 2022	DS			
	8580796*PED	Mar 25, 2023				
	8642600	Jan 28, 2022		U-1492		
	8642600*PED	Jul 28, 2022				
	8642760	Sep 25, 2022	DS			
	8642760*PED	Mar 25, 2023				
	8759350	Mar 02, 2027		U-1529		
	9089567	Jan 28, 2022		U-543		
	9125939	Jul 28, 2026		U-1749		
	9358207	Apr 12, 2020	DP			
	9359302	Sep 25, 2022	DS DP	U-1859		

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APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>ARIPIRAZOLE - ABILIFY</u>						
N 021729	003	9387182				
		Dec 25, 2023	U-1529			
<u>ARIPIRAZOLE - ABILIFY</u>						
N 021729	004	7053092				
		Jan 28, 2022	U-839		ODE-80	Dec 12, 2021
		8017615		DP		
		8017615*PED				
		Dec 16, 2024				
		8518421		DP		
		8518421*PED				
		Jul 24, 2021				
		8580796		DS		
		8580796*PED				
		Mar 25, 2023				
		8642600			U-1492	
		8642600*PED				
		Jul 28, 2022				
		8642760		DS		
		8642760*PED				
		Mar 25, 2023				
		9358207		DP		
		9359302		DS DP	U-1859	
		Sep 25, 2022				
		9387182			U-1529	
		Dec 25, 2023				
<u>ARIPIRAZOLE - ABILIFY</u>						
N 021729	005	7053092				
		Jan 28, 2022	U-839		ODE-80	Dec 12, 2021
		8017615		DP		
		8017615*PED				
		Dec 16, 2024				
		8518421		DP		
		8518421*PED				
		Jul 24, 2021				
		8580796		DS		
		8580796*PED				
		Mar 25, 2023				
		8642600			U-1492	
		8642600*PED				
		Jul 28, 2022				
		8642760		DS		
		8642760*PED				
		Mar 25, 2023				
		9358207		DP		
		9359302		DS DP	U-1859	
		Sep 25, 2022				
		9387182			U-1529	
		Dec 25, 2023				
<u>ARIPIRAZOLE - ABILIFY</u>						
N 021866	001	7115587				
		Jul 21, 2024	DP U-764		ODE-80	Dec 12, 2021
		7115587*PED				
		Jan 21, 2025				
		7550445		DP		
		Jul 21, 2024				
<u>ARIPIRAZOLE - ABILIFY MAINTENA KIT</u>						
N 202971	001	10525057				
		Mar 08, 2034	U-1632		I-746	Jul 27, 2020
		10525057				
		Mar 08, 2034	U-2723			
		10525057				
		Mar 08, 2034	U-543			
		7807680		DP		
		Oct 19, 2024				
		8030313			U-1632	
		Oct 19, 2024				
		8030313			U-543	
		Oct 19, 2024				
		8338427		DP	U-1633	
		Mar 15, 2025				
		8338427		DP	U-543	
		Mar 15, 2025				
		8338428		DP	U-1633	
		Aug 06, 2023				
		8338428		DP	U-543	
		Aug 06, 2023				
		8399469		DS		
		Jun 29, 2025				
		8722679		DP		
		Oct 19, 2024				
		8759351		DP	U-1530	
		Aug 06, 2023				
		8759351		DP	U-1633	
		Aug 06, 2023				
		8993761		DS		
		Sep 25, 2022				
		9089567			U-543	
		Jan 28, 2022				
<u>ARIPIRAZOLE - ABILIFY MAINTENA KIT</u>						
N 202971	002	10525057				
		Mar 08, 2034	U-1632		I-746	Jul 27, 2020
		10525057				
		Mar 08, 2034	U-2723			
		10525057				
		Mar 08, 2034	U-543			
		7807680		DP		
		Oct 19, 2024				
		8030313			U-1632	
		Oct 19, 2024				
		8030313			U-543	
		Oct 19, 2024				
		8338427		DP	U-1633	
		Mar 15, 2025				
		8338427		DP	U-543	
		Mar 15, 2025				
		8338428		DP	U-1633	
		Aug 06, 2023				
		8338428		DP	U-543	
		Aug 06, 2023				
		8399469		DS		
		Jun 29, 2025				
		8722679		DP		
		Oct 19, 2024				

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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 002	8759351	Aug 06, 2023	DP U-1530			
	8759351	Aug 06, 2023	DP U-1633			
	8993761	Sep 25, 2022	DS			
	9089567	Jan 28, 2022	U-543			
<u>ARIPIRAZOLE - ABILIFY MAINTENA KIT</u>						
N 202971 003	10525057	Mar 08, 2034	U-1632		I-746	Jul 27, 2020
	10525057	Mar 08, 2034	U-2723			
	10525057	Mar 08, 2034	U-543			
	7807680	Oct 19, 2024	DP			
	8030313	Oct 19, 2024	U-1632			
	8030313	Oct 19, 2024	U-543			
	8338427	Mar 15, 2025	DP U-1633			
	8338427	Mar 15, 2025	DP U-543			
	8338428	Aug 06, 2023	DP U-1633			
	8338428	Aug 06, 2023	DP U-543			
	8399469	Jun 29, 2025	DS			
	8722679	Oct 19, 2024	DP			
	8759351	Aug 06, 2023	DP U-1530			
	8759351	Aug 06, 2023	DP U-1633			
	8993761	Sep 25, 2022	DS			
	9089567	Jan 28, 2022	U-543			
<u>ARIPIRAZOLE - ABILIFY MAINTENA KIT</u>						
N 202971 004	10525057	Mar 08, 2034	U-1632		I-746	Jul 27, 2020
	10525057	Mar 08, 2034	U-2723			
	10525057	Mar 08, 2034	U-543			
	7807680	Oct 19, 2024	DP			
	8030313	Oct 19, 2024	U-1632			
	8030313	Oct 19, 2024	U-543			
	8338427	Mar 15, 2025	DP U-1633			
	8338427	Mar 15, 2025	DP U-543			
	8338428	Aug 06, 2023	DP U-1633			
	8338428	Aug 06, 2023	DP U-543			
	8399469	Jun 29, 2025	DS			
	8722679	Oct 19, 2024	DP			
	8759351	Aug 06, 2023	DP U-1530			
	8759351	Aug 06, 2023	DP U-1633			
	8993761	Sep 25, 2022	DS			
	9089567	Jan 28, 2022	U-543			
<u>ARIPIRAZOLE - ABILIFY MYCITE KIT</u>						
N 207202 001	10441194	May 12, 2029	DP		I-746	Jul 27, 2020
	10517507	Jun 13, 2032	DP			
	7053092	Jan 28, 2022	U-1529			
	7978064	Sep 14, 2026	DP			
	8017615	Jun 16, 2024	DP			
	8114021	Jun 21, 2030	DP			
	8258962	Nov 25, 2030	DP			
	8545402	Apr 27, 2030	DP			
	8547248	Dec 18, 2030	DP U-2167			
	8580796	Sep 25, 2022	DS			
	8642760	Sep 25, 2022	DS			
	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			
	9089567	Jan 28, 2022	U-543			
	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			
	9359302	Sep 25, 2022	DS DP U-1529			
	9359302	Sep 25, 2022	DS DP U-1749			

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<u>ARIPIRAZOLE - ABILIFY MYCITE KIT</u>						
N 207202 001	9359302	Sep 25, 2022	DS DP	U-543		
	9387182	Dec 25, 2023		U-1529		
	9433371	Sep 15, 2029	DP			
	9444503	Nov 19, 2027	DP	U-2169		
	9577864	Oct 03, 2033	DP	U-2169		
	9941931	Nov 04, 2030	DP			
<u>ARIPIRAZOLE - ABILIFY MYCITE KIT</u>						
N 207202 002	10441194	May 12, 2029	DP		I-746	Jul 27, 2020
	10517507	Jun 13, 2032	DP			
	7053092	Jan 28, 2022		U-1529		
	7978064	Sep 14, 2026	DP			
	8017615	Jun 16, 2024	DP			
	8114021	Jun 21, 2030	DP			
	8258962	Nov 25, 2030	DP			
	8545402	Apr 27, 2030	DP			
	8547248	Dec 18, 2030	DP	U-2167		
	8580796	Sep 25, 2022	DS			
	8642760	Sep 25, 2022	DS			
	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			
	9089567	Jan 28, 2022		U-543		
	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			
	9359302	Sep 25, 2022	DS DP	U-1529		
	9359302	Sep 25, 2022	DS DP	U-1749		
	9359302	Sep 25, 2022	DS DP	U-543		
	9387182	Dec 25, 2023		U-1529		
	9433371	Sep 15, 2029	DP			
	9444503	Nov 19, 2027	DP	U-2169		
	9577864	Oct 03, 2033	DP	U-2169		
	9941931	Nov 04, 2030	DP			
<u>ARIPIRAZOLE - ABILIFY MYCITE KIT</u>						
N 207202 003	10441194	May 12, 2029	DP		I-746	Jul 27, 2020
	10517507	Jun 13, 2032	DP			
	7053092	Jan 28, 2022		U-1529		
	7978064	Sep 14, 2026	DP			
	8017615	Jun 16, 2024	DP			
	8114021	Jun 21, 2030	DP			
	8258962	Nov 25, 2030	DP			
	8545402	Apr 27, 2030	DP			
	8547248	Dec 18, 2030	DP	U-2167		
	8580796	Sep 25, 2022	DS			
	8642760	Sep 25, 2022	DS			
	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			
	9089567	Jan 28, 2022		U-543		
	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		

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<u>ARIPIRAZOLE - ABILIFY MYCITE KIT</u>						
N 207202 003	9320455	Dec 15, 2031	DP			
	9359302	Sep 25, 2022	DS DP	U-1529		
	9359302	Sep 25, 2022	DS DP	U-1749		
	9359302	Sep 25, 2022	DS DP	U-543		
	9387182	Dec 25, 2023		U-1529		
	9433371	Sep 15, 2029	DP			
	9444503	Nov 19, 2027	DP	U-2169		
	9577864	Oct 03, 2033	DP	U-2169		
	9941931	Nov 04, 2030	DP			
<u>ARIPIRAZOLE - ABILIFY MYCITE KIT</u>						
N 207202 004	10441194	May 12, 2029	DP		I-746	Jul 27, 2020
	10517507	Jun 13, 2032	DP			
	7053092	Jan 28, 2022		U-1529		
	7978064	Sep 14, 2026	DP			
	8017615	Jun 16, 2024	DP			
	8114021	Jun 21, 2030	DP			
	8258962	Nov 25, 2030	DP			
	8545402	Apr 27, 2030	DP			
	8547248	Dec 18, 2030	DP	U-2167		
	8580796	Sep 25, 2022	DS			
	8642760	Sep 25, 2022	DS			
	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			
	9089567	Jan 28, 2022		U-543		
	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			
	9359302	Sep 25, 2022	DS DP	U-1529		
	9359302	Sep 25, 2022	DS DP	U-1749		
	9359302	Sep 25, 2022	DS DP	U-543		
	9387182	Dec 25, 2023		U-1529		
	9433371	Sep 15, 2029	DP			
	9444503	Nov 19, 2027	DP	U-2169		
	9577864	Oct 03, 2033	DP	U-2169		
	9941931	Nov 04, 2030	DP			
<u>ARIPIRAZOLE - ABILIFY MYCITE KIT</u>						
N 207202 005	10441194	May 12, 2029	DP		I-746	Jul 27, 2020
	10517507	Jun 13, 2032	DP			
	7053092	Jan 28, 2022		U-1529		
	7978064	Sep 14, 2026	DP			
	8017615	Jun 16, 2024	DP			
	8114021	Jun 21, 2030	DP			
	8258962	Nov 25, 2030	DP			
	8545402	Apr 27, 2030	DP			
	8547248	Dec 18, 2030	DP	U-2167		
	8580796	Sep 25, 2022	DS			
	8642760	Sep 25, 2022	DS			
	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			
	9089567	Jan 28, 2022		U-543		
	9119554	Dec 16, 2028	DP			
	9125939	Jul 28, 2026		U-1749		

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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 005	9149577	Dec 15, 2029	DP			
	9258035	Mar 05, 2029	DP			
	9268909	Oct 15, 2033	DP	U-2168		
	9320455	Dec 15, 2031	DP			
	9359302	Sep 25, 2022	DS DP	U-1529		
	9359302	Sep 25, 2022	DS DP	U-1749		
	9359302	Sep 25, 2022	DS DP	U-543		
	9387182	Dec 25, 2023		U-1529		
	9433371	Sep 15, 2029	DP			
	9444503	Nov 19, 2027	DP	U-2169		
	9577864	Oct 03, 2033	DP	U-2169		
	9941931	Nov 04, 2030	DP			
<u>ARIPIRAZOLE - ABILIFY MYCITE KIT</u>						
N 207202 006	10441194	May 12, 2029	DP		I-746	Jul 27, 2020
	10517507	Jun 13, 2032	DP			
	7053092	Jan 28, 2022		U-1529		
	7978064	Sep 14, 2026	DP			
	8017615	Jun 16, 2024	DP			
	8114021	Jun 21, 2030	DP			
	8258962	Nov 25, 2030	DP			
	8545402	Apr 27, 2030	DP			
	8547248	Dec 18, 2030	DP	U-2167		
	8580796	Sep 25, 2022	DS			
	8642760	Sep 25, 2022	DS			
	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			
	9089567	Jan 28, 2022		U-543		
	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			
	9359302	Sep 25, 2022	DS DP	U-1529		
	9359302	Sep 25, 2022	DS DP	U-1749		
	9359302	Sep 25, 2022	DS DP	U-543		
	9387182	Dec 25, 2023		U-1529		
	9433371	Sep 15, 2029	DP			
	9444503	Nov 19, 2027	DP	U-2169		
	9577864	Oct 03, 2033	DP	U-2169		
	9941931	Nov 04, 2030	DP			
<u>ARIPIRAZOLE LAUROXIL - ARISTADA</u>						
N 207533 001	10112903	Jun 24, 2030	DS	U-543	NCE	Oct 05, 2020
	10226458	Mar 19, 2032		U-543		
	10238651	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	NCE	Oct 05, 2020
	10226458	Mar 19, 2032		U-543		
	10238651	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			

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<u>ARIPIPRAZOLE LAUROXIL - ARISTADA</u>						
N 207533 003	10112903	Jun 24, 2030	DS	U-543	NCE	Oct 05, 2020
	10226458	Mar 19, 2032		U-543		
	10238651	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>ARIPIPRAZOLE LAUROXIL - ARISTADA</u>						
N 207533 004	10112903	Jun 24, 2030	DS	U-543	NCE	Oct 05, 2020
	10226458	Mar 19, 2032		U-543		
	10238651	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>ARIPIPRAZOLE LAUROXIL - ARISTADA INITIO KIT</u>						
N 209830 001	10016415	Sep 08, 2035		DP	NCE	Oct 05, 2020
	10112903	Jun 24, 2030	DS	U-543		
	8431576	Oct 26, 2030	DS			
	8796276	Jun 24, 2030		U-543		
<u>ARMODAFINIL - NUVIGIL</u>						
N 021875 001	7132570	Dec 18, 2023	DS	DP		
	7297346	Nov 29, 2023		DP		
<u>ARMODAFINIL - NUVIGIL</u>						
N 021875 002	7132570	Dec 18, 2023	DS	DP		
	7297346	Nov 29, 2023		DP		
<u>ARMODAFINIL - NUVIGIL</u>						
N 021875 003	7132570	Dec 18, 2023	DS	DP		
	7297346	Nov 29, 2023		DP		
<u>ARMODAFINIL - NUVIGIL</u>						
N 021875 004	7132570	Dec 18, 2023	DS	DP		
	7297346	Nov 29, 2023		DP		
<u>ARMODAFINIL - NUVIGIL</u>						
N 021875 005	7132570	Dec 18, 2023	DS	DP		
	7297346	Nov 29, 2023		DP		
<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>ASCORBIC ACID - ASCOR</u>						
N 209112 001					ODE-160	Oct 02, 2024
<u>ASCORBIC ACID; POLYETHYLENE GLYCOL 3350; POTASSIUM CHLORIDE; SODIUM ASCORBATE; SODIUM CHLORIDE; SODIUM SULFATE - MOVIPREP</u>						
N 021881 001	7169381	Sep 01, 2024	DS	DP		
	7658914	Sep 01, 2024	DS	DP		
<u>ASCORBIC ACID; POLYETHYLENE GLYCOL 3350; POTASSIUM CHLORIDE; SODIUM ASCORBATE; SODIUM CHLORIDE; SODIUM SULFATE - PLENUVU</u>						
N 209381 001	10016504	Sep 10, 2033		DP	NP	May 04, 2021
	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	NP	Oct 11, 2022
	9687474	Jul 25, 2033		DP		

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<u>ASENAPINE - SECUADO</u>						
N 212268 001	10022445	Jul 25, 2033	DP		NP	Oct 11, 2022
	9687474	Jul 25, 2033	DP			
<u>ASENAPINE - SECUADO</u>						
N 212268 002	10022445	Jul 25, 2033	DP		NP	Oct 11, 2022
	9687474	Jul 25, 2033	DP			
<u>ASENAPINE - SECUADO</u>						
N 212268 003	10022445	Jul 25, 2033	DP		NP	Oct 11, 2022
	9687474	Jul 25, 2033	DP			
<u>ASENAPINE MALEATE - SAPHRIS</u>						
N 022117 001	5763476	Jun 09, 2020	DP U-1960		D-166	Jan 13, 2020
	5763476	Jun 09, 2020	DP U-1961		I-597	Jan 13, 2020
	5763476	Jun 09, 2020	DP U-1962			
	5763476	Jun 09, 2020	DP U-1963			
	5763476	Jun 09, 2020	DP U-326			
	5763476*PED	Dec 09, 2020				
	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	5763476	Jun 09, 2020	DP U-1960		D-166	Jan 13, 2020
	5763476	Jun 09, 2020	DP U-1961		I-597	Jan 13, 2020
	5763476	Jun 09, 2020	DP U-1962			
	5763476	Jun 09, 2020	DP U-1963			
	5763476	Jun 09, 2020	DP U-326			
	5763476*PED	Dec 09, 2020				
	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	5763476	Jun 09, 2020	DP U-1893		D-166	Jan 13, 2020
	5763476	Jun 09, 2020	DP U-1966		I-597	Jan 13, 2020
	5763476*PED	Dec 09, 2020				
	7741358	Apr 06, 2026	DS DP U-1893			
	7741358	Apr 06, 2026	DS DP U-1966			
	7741358*PED	Oct 06, 2026				
	8022228	Apr 06, 2026	DS DP			
	8022228*PED	Oct 06, 2026				
<u>ASPIRIN - VAZALORE</u>						
N 203697 001	8865187	Mar 23, 2022	DP			
	9101637	Mar 23, 2022		U-1731		
	9101637	Mar 23, 2022		U-1732		
	9101637	Mar 23, 2022		U-1733		
	9216150	Sep 29, 2032	DP			
	9226892	Sep 29, 2032		U-1731		
	9226892	Sep 29, 2032		U-1732		
	9226892	Sep 29, 2032		U-1733		
	9351984	Dec 19, 2021	DP			
<u>ASPIRIN; OMEPRAZOLE - YOSPRALA</u>						
N 205103 001	6926907	Feb 28, 2023	DP U-1902			
	8206741	Feb 28, 2023	DP U-1902			
	9364439	May 31, 2022	DP U-1902			
	9539214	Mar 13, 2033		U-1902		
	9987231	Jan 02, 2033		U-2324		

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<u>ASPIRIN; OMEPRAZOLE - YOSPRALA</u>						
N 205103	001	6926907	Feb 28, 2023	DP	U-1902	
		8206741	Feb 28, 2023	DP	U-1902	
		9364439	May 31, 2022	DP	U-1902	
		9539214	Mar 13, 2033		U-1902	
		9987231	Jan 02, 2033		U-2324	
<u>ASPIRIN; OMEPRAZOLE - YOSPRALA</u>						
N 205103	002	6926907	Feb 28, 2023	DP	U-1902	
		8206741	Feb 28, 2023	DP	U-1902	
		9364439	May 31, 2022	DP	U-1902	
		9539214	Mar 13, 2033		U-1902	
		9987231	Jan 02, 2033		U-2324	
<u>ATAZANAVIR SULFATE; COBICISTAT - EVOTAZ</u>						
N 206353	001	10039718	Oct 04, 2032	DP		
		8148374	Sep 03, 2029	DS DP	U-1279	
<u>ATORVASTATIN CALCIUM - LIPITOR</u>						
N 020702	001					M-204 Jun 23, 2020
<u>ATORVASTATIN CALCIUM - LIPITOR</u>						
N 020702	002					M-204 Jun 23, 2020
<u>AVANAFIL - STENDRA</u>						
N 202276	001	6656935	Apr 27, 2025	DS DP	U-155	
		7501409	May 05, 2023		DP	
<u>AVANAFIL - STENDRA</u>						
N 202276	002	6656935	Apr 27, 2025	DS DP	U-155	
		7501409	May 05, 2023		DP	
<u>AVANAFIL - STENDRA</u>						
N 202276	003	6656935	Apr 27, 2025	DS DP	U-155	
		7501409	May 05, 2023		DP	
<u>AVAPRITINIB - AYYAKIT</u>						
N 212608	001	9200002	Oct 15, 2034	DS DP	U-2726	NCE Jan 09, 2025
		9944651	Oct 15, 2034	DS DP	U-2726	
		9994575	Oct 15, 2034	DS DP	U-2726	
<u>AVAPRITINIB - AYYAKIT</u>						
N 212608	002	9200002	Oct 15, 2034	DS DP	U-2726	NCE Jan 09, 2025
		9944651	Oct 15, 2034	DS DP	U-2726	
		9994575	Oct 15, 2034	DS DP	U-2726	
<u>AVAPRITINIB - AYYAKIT</u>						
N 212608	003	9200002	Oct 15, 2034	DS DP	U-2726	NCE Jan 09, 2025
		9944651	Oct 15, 2034	DS DP	U-2726	
		9994575	Oct 15, 2034	DS DP	U-2726	
<u>AVATROMBOPAG MALEATE - DOPTELET</u>						
N 210238	001	7638536	May 05, 2025	DS DP		I-802 Jun 26, 2022
		8338429	Jun 30, 2023		U-2577	NCE May 21, 2023
		8765764	Jan 15, 2023		U-2314	ODE-246 Jun 26, 2026
		8765764	Jan 15, 2023		U-2578	
<u>AVIBACTAM SODIUM; CEFTAZIDIME - AVYCAZ</u>						
N 206494	001	7112592	Jan 07, 2026	DS DP	U-2244	
		7112592	Jan 07, 2026	DS DP	U-2508	
		7112592	Jan 07, 2026	DS DP	U-282	
		7612087	Nov 12, 2026		DP	
		8178554	Jul 24, 2021	DS DP	U-2245	
		8178554	Jul 24, 2021	DS DP	U-2509	
		8178554	Jul 24, 2021	DS DP	U-282	
		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		

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<u>AXITINIB - INLYTA</u>						
N 202324	001	6534524	Apr 29, 2025	DS DP		
		7141581	Jun 30, 2020		U-1220	
		8791140	Dec 14, 2030	DS		
<u>AXITINIB - INLYTA</u>						
N 202324	002	6534524	Apr 29, 2025	DS DP		
		7141581	Jun 30, 2020		U-1220	
		8791140	Dec 14, 2030	DS		
<u>AZELAIC ACID - FINACEA</u>						
N 207071	001	10117812	Oct 18, 2027	DP	U-1796	
		10322085	Oct 24, 2023	DP		
		7700076	Sep 18, 2027	DP		
		8435498	Mar 01, 2024		U-1727	
		8722021	Oct 24, 2023	DP		
		8900554	Oct 24, 2023	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; FLUTICASONE PROPIONATE - AZELASTINE HYDROCHLORIDE AND FLUTICASONE PROPIONATE</u>						
A 207712	001				PC	Aug 29, 2020
<u>AZELASTINE HYDROCHLORIDE; FLUTICASONE PROPIONATE - DYMISTA</u>						
N 202236	001	8163723	Aug 29, 2023		U-1667	
		8163723	Aug 29, 2023		U-644	
		8163723	Aug 29, 2023		U-707	
		8163723	Aug 29, 2023		U-77	
		8163723	Aug 29, 2023		U-81	
		8163723*PED	Feb 29, 2024			
		8168620	Feb 24, 2026	DP		
		9259428	Jun 13, 2023		U-644	
		9259428*PED	Dec 13, 2023			
		9901585	Jun 13, 2023	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		
		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>AZITHROMYCIN - ZMAX</u>						
N 050797	001	6984403	Feb 14, 2024	DP	U-282	
		7887844	Feb 14, 2024	DP		

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<u>AZTREONAM - CAYSTON</u>						
N 050814	001	7208141	Dec 20, 2021	DP U-1031		
		7214364	Dec 20, 2021	DP		
		7427633	Dec 20, 2021	DP U-1031		
		8399496	Dec 20, 2021	DP U-1377		
<u>BALOXAVIR MARBOXIL - XOFLUZA</u>						
N 210854	001	10392406	Apr 27, 2036	DS	I-811	Oct 16, 2022
		8927710	May 05, 2031	DP	NCE	Oct 24, 2023
		8987441	Sep 21, 2031	DS DP		
		9815835	Jun 14, 2030	DP		
<u>BALOXAVIR MARBOXIL - XOFLUZA</u>						
N 210854	002	10392406	Apr 27, 2036	DS	I-811	Oct 16, 2022
		8927710	May 05, 2031	DP	NCE	Oct 24, 2023
		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		
<u>BARICITINIB - OLUMIANT</u>						
N 207924	001	8158616	Jun 08, 2030	DS DP	NCE	May 31, 2023
		8420629	Mar 10, 2029	U-247		
<u>BAZEDOXIFENE ACETATE; ESTROGENS, CONJUGATED - DUAVEE</u>						
N 022247	001	6479535	May 06, 2024	DP U-594		
		6479535	May 06, 2024	DP U-904		
		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		
		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		
		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		
<u>BECLOMETHASONE DIPROPIONATE - QVAR REDHALER</u>						
N 207921	001	10022509	May 18, 2031	DP		
		10022510	May 18, 2031	DP		
		10086156	May 18, 2031	DP		
		7637260	Aug 25, 2020	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		

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<u>BECLOMETHASONE DIPROPIONATE - QVAR REDHALER</u>						
N 207921 002	10086156	May 18, 2031	DP			
	7637260	Aug 25, 2020	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		NPP	Aug 09, 2022
	8546428	Mar 19, 2029	DS DP U-1321		ODE-251	Aug 09, 2026
<u>BELINOSTAT - BELEODAQ</u>						
N 206256 001	6888027	Sep 27, 2021	DS DP U-1544		ODE-68	Jul 03, 2021
	8835501	Oct 27, 2027	DP			
<u>BEMPEDOIC ACID - NEXLETOL</u>						
N 211616 001	10118881	Dec 23, 2023		U-2747	NCE	Feb 21, 2025
	7335799	Dec 03, 2025	DS			
	8497301	Dec 23, 2023		U-2747		
	9000041	Dec 23, 2023		U-2747		
	9624152	Dec 23, 2023		U-2748		
<u>BEMPEDOIC ACID; EZETIMIBE - NEXLIZET</u>						
N 211617 001	10118881	Dec 23, 2023		U-2746	NCE	Feb 21, 2025
	7335799	Dec 03, 2025	DS		NP	Feb 26, 2023
	8497301	Dec 23, 2023		U-2746		
	9000041	Dec 23, 2023		U-2746		
	9624152	Dec 23, 2023		U-2749		
<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			
<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				
	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			
<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				

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<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			
	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		ODE-179	Dec 07, 2022
	10052385	Mar 15, 2033	U-1971			
	10052385	Mar 15, 2033	U-1972			
	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>BENZNIDAZOLE - BENZNIDAZOLE</u>						
N 209570 001					NCE	Aug 29, 2022
					ODE-154	Aug 29, 2024
<u>BENZNIDAZOLE - BENZNIDAZOLE</u>						
N 209570 002					NCE	Aug 29, 2022
					ODE-154	Aug 29, 2024
<u>BENZOYL PEROXIDE; CLINDAMYCIN PHOSPHATE - ACANYA</u>						
N 050819 001	10220049	Jun 03, 2029	DP 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			

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<u>BENZOYL PEROXIDE; CLINDAMYCIN PHOSPHATE - ONEXTON</u>						
N 050819	002	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>BENZYL ALCOHOL - ULESFIA</u>						
N 022129	001	6793931	Jul 11, 2022	DP U-970		
		7294342	May 19, 2024	U-970		
<u>BEPOTASTINE BESILATE - BEPREVE</u>						
N 022288	001	8784789	Sep 05, 2024	DP		
		8877168	Jul 30, 2023	DP		
<u>BESIFLOXACIN HYDROCHLORIDE - BESIVANCE</u>						
N 022308	001	6685958	Jun 29, 2021	DP U-80		
		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 - TACLONEX</u>						
N 021852	001	6753013	Jan 27, 2020	DP U-193		
		6753013	Jan 27, 2020	DP U-88		
<u>BETAMETHASONE DIPROPIONATE; CALCIPOTRIENE - TACLONEX</u>						
N 022185	001	6753013	Jan 27, 2020	DP U-1761	NPP	Jul 25, 2022
		6753013	Jan 27, 2020	DP U-193	PED	Jan 25, 2023
		6753013	Jan 27, 2020	DP U-88		
		6787529	Jan 27, 2020	DP U-1761		
		6787529	Jan 27, 2020	DP U-193		
		6787529	Jan 27, 2020	DP U-88		
<u>BETAMETHASONE DIPROPIONATE; CALCIPOTRIENE - ENSTILAR</u>						
N 207589	001	10130640	Jun 10, 2031	DP	NPP	Jul 30, 2022
		10130640*PED	Dec 10, 2031		PED	Jan 30, 2023
		6753013	Jan 27, 2020	DP U-1761		
		6753013	Jan 27, 2020	DP U-2627		
		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>BETRIXABAN - BEVYXXA</u>						
N 208383	001	6376515	Sep 15, 2020	DS DP U-1167	NCE	Jun 23, 2022
		6376515	Sep 15, 2020	DS DP U-1502		
		6376515	Sep 15, 2020	DS DP U-2029		
		6376515	Sep 15, 2020	DS DP U-2030		
		6835739	Sep 15, 2020	DS DP		
		7598276	Nov 08, 2026	DS		
		8404724	Mar 29, 2031	DP U-2034		
		8518977	Sep 15, 2020	DS		
		8557852	Sep 08, 2028	U-1167		
		8557852	Sep 08, 2028	U-2030		
		8691847	Sep 15, 2020	DS DP U-2029		
		8691847	Sep 15, 2020	DS DP U-2035		

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<u>BETRIXABAN - BEVYXXA</u>						
N 208383	001	8987463	Dec 28, 2030	DP		
		9555023	Nov 07, 2026	U-1502		
		9629831	Sep 15, 2020	U-1167		
		9629831	Sep 15, 2020	U-1502		
		9629831	Sep 15, 2020	U-2030		
		9629831	Sep 15, 2020	U-2035		
<u>BETRIXABAN - BEVYXXA</u>						
N 208383	002	6376515	Sep 15, 2020	DS DP U-1167	NCE	Jun 23, 2022
		6376515	Sep 15, 2020	DS DP U-1502		
		6376515	Sep 15, 2020	DS DP U-2029		
		6376515	Sep 15, 2020	DS DP U-2030		
		6835739	Sep 15, 2020	DS DP		
		7598276	Nov 08, 2026	DS		
		8404724	Mar 29, 2031	DP U-2034		
		8518977	Sep 15, 2020	DS		
		8557852	Sep 08, 2028	U-1167		
		8557852	Sep 08, 2028	U-2030		
		8691847	Sep 15, 2020	DS DP U-2029		
		8691847	Sep 15, 2020	DS DP U-2035		
		8987463	Dec 28, 2030	DP		
		9555023	Nov 07, 2026	U-1502		
		9629831	Sep 15, 2020	U-1167		
		9629831	Sep 15, 2020	U-1502		
		9629831	Sep 15, 2020	U-2030		
		9629831	Sep 15, 2020	U-2035		
<u>BICTEGRAVIR SODIUM; EMTRICITABINE; TENOFOVIR ALAFENAMIDE FUMARATE - BIKTARVY</u>						
N 210251	001	10385067	Jun 19, 2035	U-257	NCE	Feb 07, 2023
		10548846	Nov 08, 2036	DP	NPP	Jun 18, 2022
		6642245	Nov 04, 2020	U-257	ODE-256	Jun 18, 2026
		6703396	Mar 09, 2021	DS DP		
		7390791	May 07, 2022	DS DP		
		7803788	Feb 02, 2022	U-257		
		8754065	Aug 15, 2032	DS DP U-257		
		9216996	Dec 19, 2033	DS DP		
		9296769	Aug 15, 2032	DS DP U-257		
		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 - LATISSE</u>						
N 022369	001	8038988	Aug 25, 2023	DS DP U-1208		
		8101161	May 25, 2024	U-1217		
		8101161	May 25, 2024	U-1218		
		8263054	Jan 15, 2023	U-1277		
		8541466	Jan 31, 2021	U-1217		
		8632760	Jan 15, 2023	U-1487		
		8758733	Jan 15, 2023	U-1487		
		8906962	Jan 31, 2021	U-1217		
		8986715	Jan 15, 2023	U-1217		
		9216183	Jan 15, 2023	U-1487		
		9226931	Jan 15, 2023	U-1799		
		9579270	Jan 31, 2021	U-1975		

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<u>BINIMETINIB - MEKTOVI</u>						
N 210498 001	10005761	Aug 27, 2030		U-2331	NCE	Jun 27, 2023
	7777050	Mar 13, 2023	DS DP		ODE-194	Jun 27, 2025
	8178693	Mar 13, 2023	DS DP			
	8193229	Mar 13, 2023		U-2330		
	8513293	Mar 13, 2023		U-2331		
	9314464	Jul 04, 2031		U-2332		
	9562016	Oct 18, 2033	DS DP			
	9593100	Aug 27, 2030	DP			
	9598376	Oct 18, 2033		U-2330		
	9850229	Aug 27, 2030		U-2333		
	9980944	Oct 18, 2033		U-2334		
<u>BISACODYL; POLYETHYLENE GLYCOL 3350; POTASSIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE - HALFLYTELY</u>						
N 021551 003	7291324	Oct 22, 2022		U-837		
<u>BIVALIRUDIN - ANGIOMAX</u>						
N 020873 001	7582727	Jul 27, 2028		DP		
	7598343	Jul 27, 2028		DP		
<u>BOCEPREVIR - VICTRELIS</u>						
N 202258 001	7772178	Nov 11, 2027		DP U-1128		
	8119602	Mar 17, 2027		U-1233		
	RE43298	Dec 22, 2024	DS DP	U-1128		
<u>BORTEZOMIB - VELCADE</u>						
N 021602 001	6713446	Jan 25, 2022	DS DP		ODE-76	Oct 08, 2021
	6713446*PED	Jul 25, 2022			PED	Apr 08, 2022
	6958319	Jan 25, 2022	DS DP			
	6958319*PED	Jul 25, 2022				
<u>BORTEZOMIB - BORTEZOMIB</u>						
N 205004 001	8962572	Nov 03, 2032		DP		
<u>BOSENTAN - TRACLEER</u>						
N 021290 001					NPP	Sep 05, 2020
<u>BOSENTAN - TRACLEER</u>						
N 021290 002					NPP	Sep 05, 2020
<u>BOSENTAN - TRACLEER</u>						
N 209279 001	7959945	Dec 28, 2027		DP	NP	Sep 05, 2020
	8309126	May 15, 2026		DP	ODE-161	Sep 05, 2024
<u>BOSUTINIB MONOHYDRATE - BOSULIF</u>						
N 203341 001	7417148	Dec 11, 2025		U-1283	I-759	Dec 19, 2020
	7767678	Nov 23, 2026	DS DP		ODE-163	Dec 19, 2024
	7919625	Dec 11, 2025		DP		
	RE42376	Apr 13, 2024	DS			
<u>BOSUTINIB MONOHYDRATE - BOSULIF</u>						
N 203341 002	7417148	Dec 11, 2025		U-1283	I-759	Dec 19, 2020
	7767678	Nov 23, 2026	DS DP		ODE-163	Dec 19, 2024
	7919625	Dec 11, 2025		DP		
	RE42376	Apr 13, 2024	DS			
<u>BOSUTINIB MONOHYDRATE - BOSULIF</u>						
N 203341 003	7417148	Dec 11, 2025		U-1283	I-759	Dec 19, 2020
	7767678	Nov 23, 2026	DS DP		ODE-163	Dec 19, 2024
	7919625	Dec 11, 2025		DP		
	RE42376	Apr 13, 2024	DS			
<u>BREMELANOTIDE ACETATE - VYLEESI (AUTOINJECTOR)</u>						
N 210557 001	10286034	Nov 05, 2033		U-2568	NCE	Jun 21, 2024
	6579968	Jun 28, 2020	DS DP			
	6794489	Jun 28, 2020	DS DP			
	9352013	Nov 05, 2033		U-2568		
	9700592	Nov 05, 2033		U-2568		

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<u>BREXANOLONE - ZULRESSO</u>						
N 211371 001	10117951	Mar 13, 2029	DP		NCE	Jun 17, 2024
	10251894	Nov 27, 2033	U-2552			
	10322139	Jan 23, 2033	DP			
	7635773	Mar 13, 2029	DP			
	8410077	Mar 13, 2029	DP			
	9200088	Mar 13, 2029	DP			
	9750822	Mar 13, 2029	DP			
<u>BREXPIRAZOLE - REXULTI</u>						
N 205422 001	10307419	Oct 12, 2032	DP		NCE	Jul 10, 2020
	7888362	Apr 12, 2026	DS			
	8349840	Apr 12, 2026	DP U-1529			
	8618109	Apr 12, 2026	U-543			
	9839637	Apr 12, 2026	DP U-1529			
	9839637	Apr 12, 2026	DP U-543			
<u>BREXPIRAZOLE - REXULTI</u>						
N 205422 002	10307419	Oct 12, 2032	DP		NCE	Jul 10, 2020
	7888362	Apr 12, 2026	DS			
	8349840	Apr 12, 2026	DP U-1529			
	8618109	Apr 12, 2026	U-543			
	9839637	Apr 12, 2026	DP U-1529			
	9839637	Apr 12, 2026	DP U-543			
<u>BREXPIRAZOLE - REXULTI</u>						
N 205422 003	10307419	Oct 12, 2032	DP		NCE	Jul 10, 2020
	7888362	Apr 12, 2026	DS			
	8349840	Apr 12, 2026	DP U-1529			
	8618109	Apr 12, 2026	U-543			
	9839637	Apr 12, 2026	DP U-1529			
	9839637	Apr 12, 2026	DP U-543			
<u>BREXPIRAZOLE - REXULTI</u>						
N 205422 004	10307419	Oct 12, 2032	DP		NCE	Jul 10, 2020
	7888362	Apr 12, 2026	DS			
	8349840	Apr 12, 2026	DP U-1529			
	8618109	Apr 12, 2026	U-543			
	9839637	Apr 12, 2026	DP U-1529			
	9839637	Apr 12, 2026	DP U-543			
<u>BREXPIRAZOLE - REXULTI</u>						
N 205422 005	10307419	Oct 12, 2032	DP		NCE	Jul 10, 2020
	7888362	Apr 12, 2026	DS			
	8349840	Apr 12, 2026	DP U-1529			
	8618109	Apr 12, 2026	U-543			
	9839637	Apr 12, 2026	DP U-1529			
	9839637	Apr 12, 2026	DP U-543			
<u>BREXPIRAZOLE - REXULTI</u>						
N 205422 006	7888362	Apr 12, 2026	DS		NCE	Jul 10, 2020
	8349840	Apr 12, 2026	DP U-1529			
	8618109	Apr 12, 2026	U-543			
	9839637	Apr 12, 2026	DP U-1529			
	9839637	Apr 12, 2026	DP U-543			
<u>BRIGATINIB - ALUNBRIG</u>						
N 208772 001	10385078	Nov 10, 2035	DS DP U-1927		NCE	Apr 28, 2022
	9012462	Jul 31, 2030	DS		ODE-142	Apr 28, 2024
	9273077	May 21, 2029	U-1927			
	9611283	Apr 10, 2034	U-1927			
<u>BRIGATINIB - ALUNBRIG</u>						
N 208772 002	10385078	Nov 10, 2035	DS DP U-1927		NCE	Apr 28, 2022
	9012462	Jul 31, 2030	DS		ODE-142	Apr 28, 2024
	9273077	May 21, 2029	U-1927			
	9611283	Apr 10, 2034	U-1927			
<u>BRIGATINIB - ALUNBRIG</u>						
N 208772 003	10385078	Nov 10, 2035	DS DP U-1927		NCE	Apr 28, 2022
	9012462	Jul 31, 2030	DS		ODE-142	Apr 28, 2024

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<u>BRIGATINIB - ALUNBRIG</u>						
N 208772	003 9273077	May 21, 2029	U-1927			
	9611283	Apr 10, 2034	U-1927			
<u>BRILLIANT BLUE G - TISSUEBLUE</u>						
N 209569	001				NCE ODE-282	Dec 20, 2024 Dec 20, 2026
<u>BRIMONIDINE TARTRATE - ALPHAGAN P</u>						
N 021262	001 6562873	Jul 10, 2021				
	6627210	Jul 18, 2021	DP			
	6641834	Jul 28, 2021	DP			
	6673337	Jul 26, 2021	DP			
	9295641	Jul 10, 2021	U-1833			
	9295641*PED	Jan 10, 2022				
<u>BRIMONIDINE TARTRATE - OOLIANA</u>						
N 021764	001 7265117	Aug 19, 2025	DP			
<u>BRIMONIDINE TARTRATE - ALPHAGAN P</u>						
N 021770	001 10307368	Jul 10, 2021	DP			
	6562873	Jul 10, 2021	DP			
	6627210	Jul 18, 2021	DP			
	6641834	Jul 28, 2021	DP			
	6673337	Jul 26, 2021	DP			
	8858961	Sep 02, 2023	DP			
	8858961*PED	Mar 02, 2024				
	9295641	Jul 10, 2021	U-1833			
	9295641*PED	Jan 10, 2022				
	9687443	Jul 10, 2021	DP			
	9687443*PED	Jan 10, 2022				
<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			
	8859551	May 25, 2024	U-1428			
	9861631	Mar 25, 2031	U-1428			
	9861632	Mar 25, 2031	U-1428			
<u>BRIMONIDINE TARTRATE - LUMIFY</u>						
N 208144	001 8293742	Jul 14, 2030	U-2222		NP	Dec 22, 2020
<u>BRIMONIDINE TARTRATE; BRINZOLAMIDE - SIMBRINZA</u>						
N 204251	001 9044484	Oct 30, 2030	DP			
	9421265	Jun 17, 2030	DP			
<u>BRIMONIDINE TARTRATE; TIMOLOL MALEATE - COMBIGAN</u>						
N 021398	001 7030149	Apr 19, 2022	U-849			
	7320976	Apr 19, 2022	U-849			
	7323463	Jan 19, 2023	DP		Y	
	7642258	Apr 19, 2022	DS DP U-1024			
	8133890	Apr 19, 2022	U-1235			
	8354409	Apr 19, 2022	DP U-1371			
	8748425	Apr 19, 2022	DP U-1524			
	9474751	Apr 19, 2022	DP U-1524			
	9770453	Apr 19, 2022	DP U-2131			
	9907801	Apr 19, 2022	DP U-2239			
	9907802	Apr 19, 2022	DP U-2240			
<u>BRIVARACETAM - BRIVIACT</u>						
N 205836	001 6784197	Feb 21, 2021	DS DP U-2295		NCE	May 12, 2021
	6911461	Feb 21, 2021	DS DP U-2295			
	8492416	Feb 21, 2021	U-2295			

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<u>BRIVARACETAM - BRIVIACT</u>						
N 205836 002	6784197	Feb 21, 2021	DS DP U-2295		NCE	May 12, 2021
	6911461	Feb 21, 2021	DS DP U-2295			
	8492416	Feb 21, 2021	U-2295			
<u>BRIVARACETAM - BRIVIACT</u>						
N 205836 003	6784197	Feb 21, 2021	DS DP U-2295		NCE	May 12, 2021
	6911461	Feb 21, 2021	DS DP U-2295			
	8492416	Feb 21, 2021	U-2295			
<u>BRIVARACETAM - BRIVIACT</u>						
N 205836 004	6784197	Feb 21, 2021	DS DP U-2295		NCE	May 12, 2021
	6911461	Feb 21, 2021	DS DP U-2295			
	8492416	Feb 21, 2021	U-2295			
<u>BRIVARACETAM - BRIVIACT</u>						
N 205836 005	6784197	Feb 21, 2021	DS DP U-2295		NCE	May 12, 2021
	6911461	Feb 21, 2021	DS DP U-2295			
	8492416	Feb 21, 2021	U-2295			
<u>BRIVARACETAM - BRIVIACT</u>						
N 205837 001	6784197	Feb 21, 2021	DS DP U-1815		NCE	May 12, 2021
	6784197	Feb 21, 2021	DS DP U-2130			
	6911461	Feb 21, 2021	DS DP U-1815			
	6911461	Feb 21, 2021	DS DP U-2130			
	8492416	Feb 21, 2021	U-1815			
	8492416	Feb 21, 2021	U-2130			
<u>BRIVARACETAM - BRIVIACT</u>						
N 205838 001	6784197	Feb 21, 2021	DS DP U-2295		NCE	May 12, 2021
	6911461	Feb 21, 2021	DS DP U-2295			
	8492416	Feb 21, 2021	U-2295			
<u>BROMFENAC SODIUM - PROLENSA</u>						
N 203168 001	10085958	Nov 19, 2032	DP			
	8129431	Sep 11, 2025	DS DP			
	8669290	Jan 16, 2024	DP			
	8754131	Jan 16, 2024	DP			
	8871813	Jan 16, 2024	DP			
	8927606	Jan 16, 2024	U-100			
	8927606	Jan 16, 2024	U-1095			
	8927606	Jan 16, 2024	U-810			
	9144609	Jan 16, 2024	DP			
	9517220	Nov 11, 2033	U-1933			
	9561277	Jan 16, 2024	U-1933			
<u>BROMFENAC SODIUM - BROMSITE</u>						
N 206911 001	8778999	Aug 07, 2029	DP U-1834			
<u>BROMOCRIPTINE MESYLATE - CYCLOSET</u>						
N 020866 001	7888310	Jul 25, 2023	U-1433			
	8137992	Jul 25, 2023	U-1433			
	8137993	Jul 25, 2023	U-1433			
	8137994	Jul 25, 2023	U-1433			
	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			

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<u>BROMOCRIPTINE MESYLATE - CYCLOSET</u>						
N 020866 001	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			
	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	10064878	Jun 09, 2020	DP U-1325			
	10105374	Jun 09, 2020	DP U-1325			
	10143698	Jun 09, 2020	DP U-1325			
	10307375	Sep 07, 2031	DP			
	7410651	Jun 09, 2020	DP U-1325			
	7431943	Jun 09, 2020	DP			
	8293273	Jun 09, 2020	DP			
	8784888	Jun 09, 2020	DP			
	8895064	Sep 07, 2031	DP			
	9132093	Sep 07, 2031	DP			
	9192581	Sep 07, 2031	DP U-1325			
	9320716	Jun 09, 2020	DP U-1325			
	9532954	Jun 09, 2020	DP U-1325			
	RE43799	Jun 09, 2020	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; FORMOTEROL FUMARATE DIHYDRATE - SYMBICORT</u>						
N 021929 001	10166247	Jan 29, 2023	DP U-2001		M-210	Sep 11, 2020
	10166247	Jan 29, 2023	DP U-2002		M-214	Dec 20, 2020
	10166247	Jan 29, 2023	DP U-2122		NPP	Jan 27, 2020
	10166247*PED	Jul 29, 2023			PED	Jul 27, 2020
	7587988	Apr 10, 2026	DP			
	7587988*PED	Oct 10, 2026				
	7759328	Jan 29, 2023	DP U-2001			
	7759328	Jan 29, 2023	DP U-2002			
	7759328	Jan 29, 2023	DP U-2122			
	7759328*PED	Jul 29, 2023				
	7967011	Aug 11, 2021	DP			

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<u>BUDESONIDE; FORMOTEROL FUMARATE DIHYDRATE - SYMBICORT</u>						
N 021929 001	7967011*PED	Feb 11, 2022				
	8143239	Jan 29, 2023	DP U-2001			
	8143239	Jan 29, 2023	DP U-2002			
	8143239	Jan 29, 2023	DP U-2122			
	8143239*PED	Jul 29, 2023				
	8387615	Mar 26, 2027	DP			
	8387615*PED	Sep 26, 2027				
	8528545	Oct 16, 2028	DP			
	8528545*PED	Apr 16, 2029				
	8575137	Jan 29, 2023	DP U-2001			
	8575137	Jan 29, 2023	DP U-2002			
	8575137	Jan 29, 2023	DP U-2122			
	8575137*PED	Jul 29, 2023				
	8616196	Apr 07, 2029	DP			
	8616196*PED	Oct 07, 2029				
	8875699	Nov 10, 2024	DP			
	8875699*PED	May 10, 2025				
<u>BUDESONIDE; FORMOTEROL FUMARATE DIHYDRATE - SYMBICORT</u>						
N 021929 002	10166247	Jan 29, 2023	DP U-2001		M-210	Sep 11, 2020
	10166247	Jan 29, 2023	DP U-2002		M-214	Dec 20, 2020
	10166247	Jan 29, 2023	DP U-2122			
	10166247*PED	Jul 29, 2023				
	7587988	Apr 10, 2026	DP			
	7587988*PED	Oct 10, 2026				
	7759328	Jan 29, 2023	DP U-2001			
	7759328	Jan 29, 2023	DP U-2002			
	7759328	Jan 29, 2023	DP U-2122			
	7759328*PED	Jul 29, 2023				
	7967011	Aug 11, 2021	DP			
	7967011*PED	Feb 11, 2022				
	8143239	Jan 29, 2023	DP U-2001			
	8143239	Jan 29, 2023	DP U-2002			
	8143239	Jan 29, 2023	DP U-2122			
	8143239*PED	Jul 29, 2023				
	8387615	Mar 26, 2027	DP			
	8387615*PED	Sep 26, 2027				
	8528545	Oct 16, 2028	DP			
	8528545*PED	Apr 16, 2029				
	8575137	Jan 29, 2023	DP U-2001			
	8575137	Jan 29, 2023	DP U-2002			
	8575137	Jan 29, 2023	DP U-2122			
	8575137*PED	Jul 29, 2023				
	8616196	Apr 07, 2029	DP			
	8616196*PED	Oct 07, 2029				
	8875699	Nov 10, 2024	DP			
	8875699*PED	May 10, 2025				
<u>BUPIVACAINE - EXPAREL</u>						
N 022496 001	9585838	Dec 24, 2021	DP		I-771	Apr 06, 2021
<u>BUPIVACAINE - EXPAREL</u>						
N 022496 002	9585838	Dec 24, 2021	DP		I-771	Apr 06, 2021
<u>BUPRENORPHINE - BUTRANS</u>						
N 021306 001					M-250	Oct 13, 2020
<u>BUPRENORPHINE - BUTRANS</u>						
N 021306 002					M-250	Oct 13, 2020
<u>BUPRENORPHINE - BUTRANS</u>						
N 021306 003					M-250	Oct 13, 2020
<u>BUPRENORPHINE - BUTRANS</u>						
N 021306 004					M-250	Oct 13, 2020
<u>BUPRENORPHINE - BUTRANS</u>						
N 021306 005					M-250	Oct 13, 2020

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<u>BUPRENORPHINE - SUBLOCADE</u>						
N 209819 001	10198218	Jun 06, 2031		U-2489	NP	Nov 30, 2020
	10558394	Jun 25, 2031		DP		
	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	NP	Nov 30, 2020
	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 HYDROCHLORIDE - PROBUPHINE</u>						
N 204442 001	7736665	Apr 25, 2024		U-1878		
<u>BUPRENORPHINE HYDROCHLORIDE - BELBUCA</u>						
N 207932 001	7579019	Jan 22, 2020		U-1769		
	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	7579019	Jan 22, 2020		U-1769		
	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	7579019	Jan 22, 2020		U-1769		
	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	7579019	Jan 22, 2020		U-1769		
	8147866	Jul 23, 2027		DP U-1769		
	9655843	Jul 23, 2027		DP U-1556		
	9901539	Dec 21, 2032		U-1556		

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<u>BUPRENORPHINE HYDROCHLORIDE - BELBUCA</u>						
N 207932	005	7579019				
		8147866				
		9655843				
		9901539				
			U-1769			
			DP U-1769			
			DP U-1556			
			U-1556			
<u>BUPRENORPHINE HYDROCHLORIDE - BELBUCA</u>						
N 207932	006	7579019				
		8147866				
		9655843				
		9901539				
			U-1769			
			DP U-1769			
			DP U-1556			
			U-1556			
<u>BUPRENORPHINE HYDROCHLORIDE - BELBUCA</u>						
N 207932	007	7579019				
		8147866				
		9655843				
		9901539				
			U-1769			
			DP U-1769			
			DP U-1556			
			U-1556			
<u>BUPRENORPHINE HYDROCHLORIDE; NALOXONE HYDROCHLORIDE - SUBOXONE</u>						
N 022410	001	10285910				
		8017150				
		8475832				
		8603514				
		9687454				
		9855221				
		9931305				
			DP			
			DP			
			DP U-1411			
			DP U-1464			
			DP U-1464			
			DP			
			DP			
<u>BUPRENORPHINE HYDROCHLORIDE; NALOXONE HYDROCHLORIDE - SUBOXONE</u>						
N 022410	002	10285910				
		8017150				
		8475832				
		8603514				
		9687454				
		9855221				
		9931305				
			DP			
			DP			
			DP U-1411			
			DP U-1464			
			DP U-1464			
			DP			
			DP			
<u>BUPRENORPHINE HYDROCHLORIDE; NALOXONE HYDROCHLORIDE - SUBOXONE</u>						
N 022410	003	10285910				
		8017150				
		8475832				
		8603514				
		9687454				
		9855221				
		9931305				
			DP			
			DP			
			DP U-1411			
			DP U-1464			
			DP U-1464			
			DP			
			DP			
<u>BUPRENORPHINE HYDROCHLORIDE; NALOXONE HYDROCHLORIDE - SUBOXONE</u>						
N 022410	004	10285910				
		8017150				
		8475832				
		8603514				
		9687454				
		9855221				
		9931305				
			DP			
			DP			
			DP U-1411			
			DP U-1464			
			DP U-1464			
			DP			
			DP			
<u>BUPRENORPHINE HYDROCHLORIDE; NALOXONE HYDROCHLORIDE - ZUBSOLV</u>						
N 204242	001	8470361				
		8658198				
		8940330				
		9259421				
		9439900				
			DP U-1425			
			DP U-1494			
			DP			
			DP			
			DP			
			DP			
<u>BUPRENORPHINE HYDROCHLORIDE; NALOXONE HYDROCHLORIDE - ZUBSOLV</u>						
N 204242	002	8470361				
		8658198				
		8940330				
		9259421				
		9439900				
			DP U-1425			
			DP U-1494			
			DP			
			DP			
			DP			
			DP			
<u>BUPRENORPHINE HYDROCHLORIDE; NALOXONE HYDROCHLORIDE - ZUBSOLV</u>						
N 204242	003	8470361				
		8658198				
			DP U-1425			
			DP U-1494			

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<u>BUPRENORPHINE HYDROCHLORIDE; NALOXONE HYDROCHLORIDE - ZUBSOLV</u>						
N 204242 003	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	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 005	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	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	7579019	Jan 22, 2020	U-1521			
	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	7579019	Jan 22, 2020	U-1521			
	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	7579019	Jan 22, 2020	U-1521			
	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			
<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			

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<u>BUPROPION HYDROBROMIDE - APLENZIN</u>						
N 022108	003 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; NALTREXONE HYDROCHLORIDE - CONTRAVE</u>						
N 200063	001 10231964	Jul 02, 2034		U-1583		
	10307376	Nov 08, 2027		U-1585		
	7375111	Mar 26, 2025	DP			
	7462626	Jul 20, 2024		U-1583		
	8088786	Feb 03, 2029	DP			
	8318788	Nov 08, 2027		U-1584		
	8722085	Nov 08, 2027		U-1585		
	8815889	Jul 20, 2024		U-1586		
	8916195	Feb 02, 2030		U-1639		
	9107837	Jun 04, 2027		U-1639		
	9125868	Nov 08, 2027		U-1585		
	9248123	Jan 13, 2032		U-1808		
<u>CABAZITAXEL - JEVTANA KIT</u>						
N 201023	001 10583110	Oct 27, 2030		U-2753	M-201	May 17, 2020
	5847170	Mar 26, 2021	DS DP		M-209	Sep 14, 2020
	5847170*PED	Sep 26, 2021			PED	Nov 17, 2020
	7241907	Dec 10, 2025	DS			
	7241907*PED	Jun 10, 2026				
	8927592	Oct 27, 2030		U-1630		
	8927592*PED	Apr 27, 2031				
<u>CABOZANTINIB S-MALATE - COMETRIQ</u>						
N 203756	001 7579473	Aug 14, 2026	DS DP			
	8877776	Oct 08, 2030	DS DP	U-1617		
	9717720	Feb 10, 2032	DP			
<u>CABOZANTINIB S-MALATE - COMETRIQ</u>						
N 203756	002 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	I-760	Dec 19, 2020
	10039757	Jul 18, 2031		U-1480	I-792	Jan 14, 2022
	7579473	Aug 14, 2026	DS DP		ODE-227	Jan 14, 2026
	8497284	Sep 24, 2024		U-1220		
	8497284	Sep 24, 2024		U-1480		
	8497284	Sep 24, 2024		U-2488		
	8877776	Oct 08, 2030	DS DP			
	9724342	Jul 09, 2033	DP			
<u>CABOZANTINIB S-MALATE - CABOMETYX</u>						
N 208692	002 10034873	Jul 18, 2031		U-2488	I-760	Dec 19, 2020
	10039757	Jul 18, 2031		U-1480	I-792	Jan 14, 2022
	7579473	Aug 14, 2026	DS DP		ODE-227	Jan 14, 2026
	8497284	Sep 24, 2024		U-1220		
	8497284	Sep 24, 2024		U-1480		
	8497284	Sep 24, 2024		U-2488		
	8877776	Oct 08, 2030	DS DP			
	9724342	Jul 09, 2033	DP			
<u>CABOZANTINIB S-MALATE - CABOMETYX</u>						
N 208692	003 10034873	Jul 18, 2031		U-2488	I-760	Dec 19, 2020
	10039757	Jul 18, 2031		U-1480	I-792	Jan 14, 2022
	7579473	Aug 14, 2026	DS DP		ODE-227	Jan 14, 2026
	8497284	Sep 24, 2024		U-1220		
	8497284	Sep 24, 2024		U-1480		
	8497284	Sep 24, 2024		U-2488		
	8877776	Oct 08, 2030	DS DP			

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<u>CABOZANTINIB S-MALATE - CABOMETYX</u>						
N 208692	003 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			
	6582727	Aug 22, 2020	DP			
	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-1920		
	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		
	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>CALCITONIN SALMON RECOMBINANT - FORTICAL</u>						
N 021406	001 6440392	Feb 02, 2021	DP	U-227		
	RE40812	Feb 02, 2021	DP			
	RE43580	Feb 02, 2021	DP	U-227		
<u>CALCIUM ACETATE - PHOSLO</u>						
N 021160	002 6576665	Apr 03, 2021				
<u>CALCIUM ACETATE - PHOSLO GELCAPS</u>						
N 021160	003 6576665	Apr 03, 2021				
	6875445	Jul 30, 2021	DP			
<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 CARBONATE; FAMOTIDINE; MAGNESIUM HYDROXIDE - PEPCID COMPLETE</u>						
N 020958	001 6814978	Aug 26, 2021	DP			
<u>CALCIUM CHLORIDE; DEXTROSE; MAGNESIUM CHLORIDE; OXIGLUTATIONE; POTASSIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE; SODIUM PHOSPHATE - NAVSTEL</u>						
N 022193	001 7084130	Nov 29, 2021	DP	U-891		
<u>CALCIUM CHLORIDE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE; SODIUM PHOSPHATE - PHOXILLUM BK 4/2.5 IN PLASTIC CONTAINER</u>						
N 207026	001				ODE-85	Jan 13, 2022
<u>CALCIUM CHLORIDE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE; SODIUM PHOSPHATE - PHOXILLUM B2K 4/0 IN PLASTIC CONTAINER</u>						
N 207026	002				ODE-85	Jan 13, 2022
<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			

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<u>CANAGLIFLOZIN - INVOKANA</u>						
N 204042 001	7943582	Feb 26, 2029	DS DP U-2441		I-788	Oct 29, 2021
	7943582	Feb 26, 2029	DS DP U-2632		I-809	Sep 27, 2022
	7943582	Feb 26, 2029	DS DP U-493		M-197	Feb 01, 2020
	7943788	Jul 14, 2027	DS DP			
	8222219	Apr 11, 2025	U-2441			
	8222219	Apr 11, 2025	U-2632			
	8222219	Apr 11, 2025	U-493			
	8513202	Dec 03, 2027	DS DP U-2441			
	8513202	Dec 03, 2027	DS DP U-2632			
	8513202	Dec 03, 2027	DS DP U-493			
<u>CANAGLIFLOZIN - INVOKANA</u>						
N 204042 002	7943582	Feb 26, 2029	DS DP U-2441		I-788	Oct 29, 2021
	7943582	Feb 26, 2029	DS DP U-2632		I-809	Sep 27, 2022
	7943582	Feb 26, 2029	DS DP U-493		M-197	Feb 01, 2020
	7943788	Jul 14, 2027	DS DP			
	8222219	Apr 11, 2025	U-2441			
	8222219	Apr 11, 2025	U-2632			
	8222219	Apr 11, 2025	U-493			
	8513202	Dec 03, 2027	DS DP U-2441			
	8513202	Dec 03, 2027	DS DP U-2632			
	8513202	Dec 03, 2027	DS DP U-493			
<u>CANAGLIFLOZIN; METFORMIN HYDROCHLORIDE - INVOKAMET</u>						
N 204353 001	7943582	Feb 26, 2029	DS DP U-2441		I-788	Oct 29, 2021
	7943582	Feb 26, 2029	DS DP U-493		M-197	Feb 01, 2020
	7943788	Jul 14, 2027	DS DP			
	8222219	Apr 11, 2025	U-2441			
	8222219	Apr 11, 2025	U-493			
	8513202	Dec 03, 2027	DS DP U-2441			
	8513202	Dec 03, 2027	DS DP U-2632			
	8513202	Dec 03, 2027	DS DP U-493			
	8785403	Jul 30, 2024	DP			
<u>CANAGLIFLOZIN; METFORMIN HYDROCHLORIDE - INVOKAMET</u>						
N 204353 002	7943582	Feb 26, 2029	DS DP U-2441		I-788	Oct 29, 2021
	7943582	Feb 26, 2029	DS DP U-493		M-197	Feb 01, 2020
	7943788	Jul 14, 2027	DS DP			
	8222219	Apr 11, 2025	U-2441			
	8222219	Apr 11, 2025	U-493			
	8513202	Dec 03, 2027	DS DP U-2441			
	8513202	Dec 03, 2027	DS DP U-2632			
	8513202	Dec 03, 2027	DS DP U-493			
	8785403	Jul 30, 2024	DP			
<u>CANAGLIFLOZIN; METFORMIN HYDROCHLORIDE - INVOKAMET</u>						
N 204353 003	7943582	Feb 26, 2029	DS DP U-2441		I-788	Oct 29, 2021
	7943582	Feb 26, 2029	DS DP U-493		M-197	Feb 01, 2020
	7943788	Jul 14, 2027	DS DP			
	8222219	Apr 11, 2025	U-2441			
	8222219	Apr 11, 2025	U-493			
	8513202	Dec 03, 2027	DS DP U-2441			
	8513202	Dec 03, 2027	DS DP U-2632			
	8513202	Dec 03, 2027	DS DP U-493			
	8785403	Jul 30, 2024	DP			
<u>CANAGLIFLOZIN; METFORMIN HYDROCHLORIDE - INVOKAMET</u>						
N 204353 004	7943582	Feb 26, 2029	DS DP U-2441		I-788	Oct 29, 2021
	7943582	Feb 26, 2029	DS DP U-493		M-197	Feb 01, 2020
	7943788	Jul 14, 2027	DS DP			
	8222219	Apr 11, 2025	U-2441			
	8222219	Apr 11, 2025	U-493			
	8513202	Dec 03, 2027	DS DP U-2441			
	8513202	Dec 03, 2027	DS DP U-2632			
	8513202	Dec 03, 2027	DS DP U-493			
	8785403	Jul 30, 2024	DP			

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<u>CANAGLIFLOZIN; METFORMIN HYDROCHLORIDE - INVOKAMET XR</u>						
N 205879 001	6723340	Oct 25, 2021	DP		I-788	Oct 29, 2021
	7943582	Feb 26, 2029	DS DP U-2441			
	7943582	Feb 26, 2029	DS DP U-2632			
	7943582	Feb 26, 2029	DS DP U-493			
	7943788	Jul 14, 2027	DS DP			
	8222219	Apr 11, 2025		U-2441		
	8222219	Apr 11, 2025		U-2632		
	8222219	Apr 11, 2025		U-493		
	8513202	Dec 03, 2027	DS DP U-2441			
	8513202	Dec 03, 2027	DS DP U-2632			
	8513202	Dec 03, 2027	DS DP U-493			
	8785403	Jul 30, 2024	DP			
<u>CANAGLIFLOZIN; METFORMIN HYDROCHLORIDE - INVOKAMET XR</u>						
N 205879 002	7943582	Feb 26, 2029	DS DP U-2441		I-788	Oct 29, 2021
	7943582	Feb 26, 2029	DS DP U-2632			
	7943582	Feb 26, 2029	DS DP U-493			
	7943788	Jul 14, 2027	DS DP			
	8222219	Apr 11, 2025		U-2441		
	8222219	Apr 11, 2025		U-2632		
	8222219	Apr 11, 2025		U-493		
	8513202	Dec 03, 2027	DS DP U-2441			
	8513202	Dec 03, 2027	DS DP U-2632			
	8513202	Dec 03, 2027	DS DP U-493			
	8785403	Jul 30, 2024	DP			
<u>CANAGLIFLOZIN; METFORMIN HYDROCHLORIDE - INVOKAMET XR</u>						
N 205879 003	6723340	Oct 25, 2021	DP		I-788	Oct 29, 2021
	7943582	Feb 26, 2029	DS DP U-2441			
	7943582	Feb 26, 2029	DS DP U-2632			
	7943582	Feb 26, 2029	DS DP U-493			
	7943788	Jul 14, 2027	DS DP			
	8222219	Apr 11, 2025		U-2441		
	8222219	Apr 11, 2025		U-2632		
	8222219	Apr 11, 2025		U-493		
	8513202	Dec 03, 2027	DS DP U-2441			
	8513202	Dec 03, 2027	DS DP U-2632			
	8513202	Dec 03, 2027	DS DP U-493			
	8785403	Jul 30, 2024	DP			
<u>CANAGLIFLOZIN; METFORMIN HYDROCHLORIDE - INVOKAMET XR</u>						
N 205879 004	7943582	Feb 26, 2029	DS DP U-2441		I-788	Oct 29, 2021
	7943582	Feb 26, 2029	DS DP U-2632			
	7943582	Feb 26, 2029	DS DP U-493			
	7943788	Jul 14, 2027	DS DP			
	8222219	Apr 11, 2025		U-2441		
	8222219	Apr 11, 2025		U-2632		
	8222219	Apr 11, 2025		U-493		
	8513202	Dec 03, 2027	DS DP U-2441			
	8513202	Dec 03, 2027	DS DP U-2632			
	8513202	Dec 03, 2027	DS DP U-493			
	8785403	Jul 30, 2024	DP			
<u>CANGRELOR - KENGREAL</u>						
N 204958 001	10039780	Jul 10, 2035		U-2260	NCE	Jun 22, 2020
	6130208	Jun 29, 2023	DP	U-1715		
	8680052	Mar 09, 2033		U-1715		
	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	NCE	Sep 28, 2023
	10111840	Jun 17, 2035		U-2442	ODE-216	Sep 28, 2025
	10111840	Jun 17, 2035		U-2443		
	10137095	Jun 17, 2035		U-2454		
	10137095	Jun 17, 2035		U-2455		
	10195159	May 07, 2022	DS			

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<u>CANNABIDIOL - EPIDIOLEX</u>						
N 210365	001	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>CAPSAICIN - QUTENZA</u>						
N 022395	001	10034841	Sep 06, 2025	DP		
		10463598	Sep 05, 2023	DP		
		6239180	Jun 04, 2021	DP		
		8263059	Sep 05, 2023	U-2705		
		8821920	Mar 26, 2030	DP		
		8889113	Sep 05, 2023	U-2705		
		9226903	Dec 15, 2028	DP		
<u>CARBAMAZEPINE - EQUETRO</u>						
N 021710	001	6977253	May 19, 2024	U-693		
<u>CARBAMAZEPINE - EQUETRO</u>						
N 021710	002	6977253	May 19, 2024	U-693		
<u>CARBAMAZEPINE - EQUETRO</u>						
N 021710	003	6977253	May 19, 2024	U-693		
<u>CARBAMAZEPINE - CARNEXIV</u>						
N 206030	001	7635773	Mar 13, 2029	DP	ODE-124	Oct 07, 2023
		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; ENTACAPONE; LEVODOPA - STALEVO 50</u>						
N 021485	001	6500867	Jun 29, 2020	DP U-219		
		6797732	Jun 29, 2020	DP		
<u>CARBIDOPA; ENTACAPONE; LEVODOPA - STALEVO 100</u>						
N 021485	002	6500867	Jun 29, 2020	DP U-219		
		6797732	Jun 29, 2020	DP		
<u>CARBIDOPA; ENTACAPONE; LEVODOPA - STALEVO 150</u>						
N 021485	003	6500867	Jun 29, 2020	DP U-219		
		6797732	Jun 29, 2020	DP		
<u>CARBIDOPA; ENTACAPONE; LEVODOPA - STALEVO 200</u>						
N 021485	004	6500867	Jun 29, 2020	DP U-219		
		6797732	Jun 29, 2020	DP		
<u>CARBIDOPA; ENTACAPONE; LEVODOPA - STALEVO 75</u>						
N 021485	005	6500867	Jun 29, 2020	DP U-219		
		6797732	Jun 29, 2020	DP		
<u>CARBIDOPA; ENTACAPONE; LEVODOPA - STALEVO 125</u>						
N 021485	006	6500867	Jun 29, 2020	DP U-219		
		6797732	Jun 29, 2020	DP		
<u>CARBIDOPA; LEVODOPA - RYTARY</u>						
N 203312	001	7094427	May 29, 2022	DP U-1645	Y	
		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		

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<u>CARBIDOPA; LEVODOPA - RYTARY</u>						
N 203312 001	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	7094427	May 29, 2022	DP U-1645	Y		
	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 003	7094427	May 29, 2022	DP U-1645	Y		
	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	7094427	May 29, 2022	DP U-1645	Y		
	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 - DUOPA</u>						
N 203952 001					ODE-84	Jan 09, 2022
<u>CARBINOXAMINE MALEATE - KARBINAL ER</u>						
N 022556 001	8062667	Mar 29, 2029	DP			
	9522191	Jun 15, 2027	DP			

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<u>CARFILZOMIB - KYPROLIS</u>						
N 202714 001	7232818	Apr 14, 2025	DS DP		D-172	Sep 28, 2021
	7417042	Jul 20, 2026	DS DP			
	7491704	Apr 14, 2025		U-1260		
	7737112	Dec 07, 2027	DP			
	8129346	Apr 14, 2025		U-1260		
	8207125	Apr 14, 2025	DS DP			
	8207126	Apr 14, 2025	DP			
	8207127	Apr 14, 2025		U-1260		
	8207297	Apr 14, 2025	DS DP			
	9493582	Feb 27, 2033	DP			
	9511109	Oct 21, 2029		U-1924		
<u>CARFILZOMIB - KYPROLIS</u>						
N 202714 002	7232818	Apr 14, 2025	DS DP		D-172	Sep 28, 2021
	7417042	Jul 20, 2026	DS DP			
	7491704	Apr 14, 2025		U-1260		
	7737112	Dec 07, 2027	DP			
	8129346	Apr 14, 2025		U-1260		
	8207125	Apr 14, 2025	DS DP			
	8207126	Apr 14, 2025	DP			
	8207127	Apr 14, 2025		U-1260		
	8207297	Apr 14, 2025	DS DP			
	9493582	Feb 27, 2033	DP			
	9511109	Oct 21, 2029		U-1924		
<u>CARFILZOMIB - KYPROLIS</u>						
N 202714 003	7232818	Apr 14, 2025	DS DP		D-172	Sep 28, 2021
	7417042	Jul 20, 2026	DS DP			
	7491704	Apr 14, 2025		U-2319		
	7491704	Apr 14, 2025		U-2320		
	7737112	Dec 07, 2027	DP			
	8129346	Apr 14, 2025		U-2319		
	8129346	Apr 14, 2025		U-2320		
	8207125	Apr 14, 2025	DS DP			
	8207126	Apr 14, 2025	DP			
	8207127	Apr 14, 2025		U-2319		
	8207127	Apr 14, 2025		U-2320		
	8207297	Apr 14, 2025	DS DP			
	9493582	Feb 27, 2033	DP			
	9511109	Oct 21, 2029		U-1924		
<u>CARIPRAZINE HYDROCHLORIDE - VRAYLAR</u>						
N 204370 001	7737142	Sep 17, 2029	DS DP U-1750		I-798	May 24, 2022
	7737142	Sep 17, 2029	DS DP U-2543		M-213	Nov 09, 2020
	7737142	Sep 17, 2029	DS DP U-2544		NCE	Sep 17, 2020
	7737142	Sep 17, 2029	DS DP U-2545			
	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		
<u>CARIPRAZINE HYDROCHLORIDE - VRAYLAR</u>						
N 204370 002	7737142	Sep 17, 2029	DS DP U-1750		I-798	May 24, 2022
	7737142	Sep 17, 2029	DS DP U-2543		M-213	Nov 09, 2020
	7737142	Sep 17, 2029	DS DP U-2544		NCE	Sep 17, 2020
	7737142	Sep 17, 2029	DS DP U-2545			
	7943621	Dec 16, 2028	DS DP			
<u>CARIPRAZINE HYDROCHLORIDE - VRAYLAR</u>						
N 204370 003	7737142	Sep 17, 2029	DS DP U-1750		I-798	May 24, 2022
	7737142	Sep 17, 2029	DS DP U-2543		M-213	Nov 09, 2020
	7737142	Sep 17, 2029	DS DP U-2544		NCE	Sep 17, 2020
	7943621	Dec 16, 2028	DS DP			
<u>CARIPRAZINE HYDROCHLORIDE - VRAYLAR</u>						
N 204370 004	7737142	Sep 17, 2029	DS DP U-1750		I-798	May 24, 2022
	7737142	Sep 17, 2029	DS DP U-2543		M-213	Nov 09, 2020
	7737142	Sep 17, 2029	DS DP U-2544		NCE	Sep 17, 2020
	7943621	Dec 16, 2028	DS DP			

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<u>CARIPRAZINE HYDROCHLORIDE - VRAYLAR</u>						
N 204370	004	7737142	Sep 17, 2029	DS DP U-1750	I-798	May 24, 2022
		7737142	Sep 17, 2029	DS DP U-2543	M-213	Nov 09, 2020
		7737142	Sep 17, 2029	DS DP U-2544	NCE	Sep 17, 2020
		7943621	Dec 16, 2028	DS DP		
<u>CARVEDILOL PHOSPHATE - COREG CR</u>						
N 022012	001	7268156	Jun 27, 2023	DS DP U-3		
		7268156	Jun 27, 2023	DS DP U-313		
		8101209	Sep 11, 2025	DP		
<u>CARVEDILOL PHOSPHATE - COREG CR</u>						
N 022012	002	7268156	Jun 27, 2023	DS DP U-3		
		7268156	Jun 27, 2023	DS DP U-313		
		8101209	Sep 11, 2025	DP		
<u>CARVEDILOL PHOSPHATE - COREG CR</u>						
N 022012	003	7268156	Jun 27, 2023	DS DP U-3		
		7268156	Jun 27, 2023	DS DP U-313		
		8101209	Sep 11, 2025	DP		
<u>CARVEDILOL PHOSPHATE - COREG CR</u>						
N 022012	004	7268156	Jun 27, 2023	DS DP U-3		
		7268156	Jun 27, 2023	DS DP U-313		
		8101209	Sep 11, 2025	DP		
<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>CEFIDEROCOL SULFATE TOSYLATE - FETROJA</u>						
N 209445	001	10004750	Sep 03, 2035	DS DP	NCE	Nov 14, 2024
		9238657	Nov 19, 2031	DS DP U-282	GAIN	Nov 14, 2029
		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	6417175	Apr 11, 2022	DS DP U-1676		
		6906055	Dec 15, 2021	DS DP		
		7419973	Dec 15, 2021	DP		
		8247400	Feb 10, 2031	DP U-282		
		9629861	Sep 21, 2030	DP		
<u>CEFTAROLINE FOSAMIL - TEFLARO</u>						
N 200327	002	6417175	Apr 11, 2022	DS DP U-1676		
		6906055	Dec 15, 2021	DS DP		
		7419973	Dec 15, 2021	DP		
		8247400	Feb 10, 2031	DP U-282		
		9629861	Sep 21, 2030	DP		
<u>CEFTOLOZANE SULFATE; TAZOBACTAM SODIUM - ZERBAXA</u>						
N 206829	001	10028963	Sep 07, 2032	U-2565	NCE	Dec 19, 2019
		10028963	Sep 07, 2032	U-2566	GAIN	Dec 19, 2024
		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		
		7129232	May 15, 2028	DS DP U-1676		
		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		
		9320740	Mar 14, 2034	DP		
		9724353	Sep 07, 2032	U-2565		

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<u>Ceftolozane Sulfate; Tazobactam Sodium - Zerbaxa</u>						
N 206829 001	9724353	Sep 07, 2032				
	9872906	Mar 14, 2034	U-2566 DP			
<u>Cenobamate - Xcopri</u>						
N 212839 001	7598279	Oct 30, 2027	DS			
<u>Cenobamate - Xcopri</u>						
N 212839 002	7598279	Oct 30, 2027	DS			
<u>Cenobamate - Xcopri</u>						
N 212839 003	7598279	Oct 30, 2027	DS			
<u>Cenobamate - Xcopri</u>						
N 212839 004	7598279	Oct 30, 2027	DS			
<u>Cenobamate - Xcopri</u>						
N 212839 005	7598279	Oct 30, 2027	DS			
<u>Cenobamate - Xcopri</u>						
N 212839 006	7598279	Oct 30, 2027	DS			
<u>Ceritinib - Zykadia</u>						
N 205755 001	7153964	Feb 26, 2021	DS DP		M-199	May 26, 2020
	7893074	Apr 25, 2026	DS DP		ODE-145	May 26, 2024
	7964592	Jan 13, 2027	DS DP		ODE-66	Apr 29, 2021
	8039479	Jun 29, 2030	DS DP			
	8188276	Jan 31, 2023	DS DP			
	8377921	Nov 20, 2027			U-1179	
	8399450	Nov 20, 2027	DS DP			
	8703787	Feb 02, 2032			U-1179	
	8835430	Jan 31, 2023	DS DP			
	9018204	Jan 31, 2023	DS DP			
	9309229	Jan 18, 2032	DS DP			
	9416112	Jan 31, 2023	DS DP			
<u>Ceritinib - Zykadia</u>						
N 211225 001	7153964	Feb 26, 2021	DS DP			
	7893074	Apr 25, 2026	DS DP			
	7964592	Jan 13, 2027	DS DP			
	8039479	Jun 29, 2030	DS DP			
	8188276	Jan 31, 2023	DS DP			
	8377921	Nov 20, 2027			U-1179	
	8399450	Nov 20, 2027	DS DP			
	8703787	Feb 02, 2032			U-1179	
	8835430	Jan 31, 2023	DS DP			
	9018204	Jan 31, 2023	DS DP			
	9309229	Jan 18, 2032	DS DP			
	9416112	Jan 31, 2023	DS DP			
<u>Cetirizine Hydrochloride - Zerviate</u>						
N 208694 001	8829005	Mar 15, 2030				
	8829005*PED	Sep 15, 2030			NDF	May 30, 2020
	9254286	Jul 09, 2032	DP		PED	Nov 30, 2020
	9254286*PED	Jan 09, 2033				
	9750684	Mar 15, 2030	DP			
	9993471	Mar 15, 2030			U-1680	
<u>Cetirizine Hydrochloride - Quzyttir</u>						
N 211415 001	8263581	Feb 28, 2030				
	8314083	Feb 28, 2030			NP	Oct 04, 2022
	8513259	Feb 11, 2030			U-2635	
	9119771	Feb 11, 2030			U-2636	
	9180090	Feb 11, 2030			U-2635	
<u>Cetirizine Hydrochloride; Pseudoephedrine Hydrochloride - Zyrtec-D 12 Hour</u>						
N 021150 002	7014867	Jun 10, 2022	DP			
	7226614	Jun 10, 2022			U-295	

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APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>CHLORHEXIDINE GLUCONATE - CHLORHEXIDINE GLUCONATE</u>						
N 021669	001 7066916	Feb 17, 2024	U-737			
	7427574	Apr 25, 2026	DP			
	7595021	May 12, 2023	DP U-1022			
	7717889	Feb 27, 2025	DP U-1022			
	7935093	Oct 02, 2027	DP U-1022			
<u>CHLORHEXIDINE GLUCONATE - READYPREP CHG</u>						
N 207964	001				NP	Nov 20, 2021
<u>CHLORHEXIDINE GLUCONATE; ISOPROPYL ALCOHOL - CHLORAPREP ONE-STEP</u>						
N 020832	001 6536975	Nov 10, 2020	DP			
<u>CHLORHEXIDINE GLUCONATE; ISOPROPYL ALCOHOL - CHLORAPREP WITH TINT</u>						
N 020832	002 6729786	Mar 14, 2023	DP			
	6991394	Jan 31, 2024	DP			
	7182536	Dec 30, 2023	DP			
	7241065	Mar 14, 2023	DP			
	7422388	Apr 25, 2027	DP U-1397			
<u>CHLORHEXIDINE GLUCONATE; ISOPROPYL ALCOHOL - CHLORAPREP ONE-STEP</u>						
N 020832	004 6536975	Nov 10, 2020	DP			
<u>CHLORHEXIDINE GLUCONATE; ISOPROPYL ALCOHOL - CHLORAPREP WITH TINT</u>						
N 020832	005 6536975	Nov 10, 2020	DP			
	6729786	Mar 14, 2023	DP			
	7241065	Mar 14, 2023	DP			
	7422388	Apr 25, 2027	DP U-1397			
<u>CHLORHEXIDINE GLUCONATE; ISOPROPYL ALCOHOL - CHLORAPREP ONE-STEP</u>						
N 020832	006 6991394	Jan 31, 2024	DP			
	7182536	Dec 30, 2023	DP			
<u>CHLORHEXIDINE GLUCONATE; ISOPROPYL ALCOHOL - CHLORAPREP WITH TINT</u>						
N 020832	007 6536975	Nov 10, 2020	DP			
	6729786	Mar 14, 2023	DP			
	7241065	Mar 14, 2023	DP			
	7422388	Apr 25, 2027	DP U-1397			
<u>CHLORHEXIDINE GLUCONATE; ISOPROPYL ALCOHOL - PREVANTICS SWAB</u>						
N 021524	001				M-221	Feb 14, 2021
<u>CHLORHEXIDINE GLUCONATE; ISOPROPYL ALCOHOL - PREVANTICS SWABSTICK</u>						
N 021524	002				M-221	Feb 14, 2021
<u>CHLORHEXIDINE GLUCONATE; ISOPROPYL ALCOHOL - PREVANTICS MAXI SWABSTICK</u>						
N 021524	003				M-221	Feb 14, 2021
<u>CHLORHEXIDINE GLUCONATE; ISOPROPYL ALCOHOL - SOLUPREP</u>						
N 208288	001 8623935	Jul 26, 2029	DP U-1022		NP	Aug 08, 2021
<u>CHLOROPROCAINE HYDROCHLORIDE - CLOROTEKAL</u>						
N 208791	001 8969412	Sep 05, 2026	DP U-2609		NP	Sep 26, 2020
	9504666	Dec 11, 2033	DP			
<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 MALEATE; IBUPROFEN; PSEUDOEPHEDRINE HYDROCHLORIDE - CHILDREN'S ADVIL ALLERGY SINUS</u>						
N 021587	001 10238640	May 25, 2024	DP			
<u>CHLORPHENIRAMINE POLISTIREX; CODEINE POLISTIREX - TUZISTRA XR</u>						
N 207768	001 8062667	Mar 29, 2029	DP			
	8790700	Mar 15, 2027	DP			
<u>CHLORZOXAZONE - CHLORZOXAZONE</u>						
A 212253	001				CGT	May 25, 2020

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<u>CHLORZOXAZONE - CHLORZOXAZONE</u>						
A 212253	002				CGT	May 25, 2020
<u>CHOLIC ACID - CHOLBAM</u>						
N 205750	001				NCE ODE-91	Mar 17, 2020 Mar 17, 2022
<u>CHOLIC ACID - CHOLBAM</u>						
N 205750	002				NCE ODE-91	Mar 17, 2020 Mar 17, 2022
<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>CHORIOGONADOTROPIN ALFA - OVIDREL</u>						
N 021149	002	6706681	Mar 16, 2021	DP		
<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	6767901	Oct 21, 2020	DP		
		8371292	Feb 01, 2028		U-1356	
		8383611	Oct 20, 2020	DP		
<u>CICLESONIDE - ZETONNA</u>						
N 202129	001	8371292	Feb 01, 2028		U-1357	
<u>CILASTATIN SODIUM; IMPENEM; RELEBACTAM - RECARBRIO</u>						
N 212819	001	8487093	Nov 19, 2029	DS DP	U-2586	
		8487093	Nov 19, 2029	DS DP	U-2587	
<u>CINACALCET HYDROCHLORIDE - SENSIPAR</u>						
N 021688	001	7829595	Sep 22, 2026	DP	U-1098	M-200 May 23, 2020
		9375405	Sep 22, 2026	DP		ODE-78 Nov 21, 2021
<u>CINACALCET HYDROCHLORIDE - SENSIPAR</u>						
N 021688	002	7829595	Sep 22, 2026	DP	U-1098	M-200 May 23, 2020
		9375405	Sep 22, 2026	DP		ODE-78 Nov 21, 2021
<u>CINACALCET HYDROCHLORIDE - SENSIPAR</u>						
N 021688	003	7829595	Sep 22, 2026	DP	U-1098	M-200 May 23, 2020
		9375405	Sep 22, 2026	DP		ODE-78 Nov 21, 2021
<u>CIPROFLOXACIN - OTIPRIO</u>						
N 207986	001	8318817	Apr 27, 2030		U-1792	I-770 Mar 02, 2021
		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 - PROQUIN XR</u>						
N 021744	001	6488962	Jun 20, 2020		DP	
<u>CIPROFLOXACIN HYDROCHLORIDE; FLUOCINOLONE ACETONIDE - OTOVEL</u>						
N 208251	001	8932610	Mar 24, 2030		DP	U-1578
<u>CIPROFLOXACIN; CIPROFLOXACIN HYDROCHLORIDE - CIPRO XR</u>						
N 021473	001	7709022	Jun 23, 2021		DP	
		8187632	Jun 23, 2021		DP	
		8187632*PED	Dec 23, 2021			
<u>CIPROFLOXACIN; CIPROFLOXACIN HYDROCHLORIDE - CIPRO XR</u>						
N 021473	002	7709022	Jun 23, 2021		DP	
		8187632	Jun 23, 2021		DP	
		8187632*PED	Dec 23, 2021			

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<u>CIPROFLOXACIN; CIPROFLOXACIN HYDROCHLORIDE - CIPRO XR</u>						
N 021473	002 7709022	Jun 23, 2021	DP			
	8187632	Jun 23, 2021	DP			
	8187632*PED	Dec 23, 2021				
<u>CIPROFLOXACIN; DEXAMETHASONE - CIPRODEX</u>						
N 021537	001 6284804	Aug 10, 2020				
	6359016	Aug 10, 2020				
	8846650	Jun 04, 2025	DP U-1578			
	9149486	Sep 13, 2022	DP U-1578			
	9345714	Sep 13, 2022	DP U-1578			
	9402805	Sep 13, 2022	DP U-1578			
	9402805	Sep 13, 2022	DP U-1679			
<u>CITRIC ACID; MAGNESIUM OXIDE; SODIUM PICOSULFATE - PREPOPIK</u>						
N 202535	001 8450338	Oct 10, 2028	DP		NPP	Aug 15, 2021
	8481083	Oct 10, 2028	DP			
<u>CITRIC ACID; MAGNESIUM OXIDE; SODIUM PICOSULFATE - CLENPIO</u>						
N 209589	001 9827231	Jun 26, 2034	DP U-2162			
<u>CLADRIBINE - MAVENCLAD</u>						
N 022561	001 7713947	Oct 16, 2026	U-2520		NP	Mar 29, 2022
	7888328	Apr 11, 2024	DP U-2521			
	8377903	May 31, 2026	U-2522			
	8785415	Apr 11, 2024	DP U-2523			
<u>CLEVIDIPINE - CLEVIPREX</u>						
N 022156	001 10010537	Oct 10, 2031	DP			
	5856346	Jan 05, 2021	DS DP U-893			
	8658676	Oct 10, 2031	DP			
<u>CLEVIDIPINE - CLEVIPREX</u>						
N 022156	002 10010537	Oct 10, 2031	DP			
	5856346	Jan 05, 2021	DS DP U-893			
	8658676	Oct 10, 2031	DP			
<u>CLEVIDIPINE - CLEVIPREX</u>						
N 022156	003 10010537	Oct 10, 2031	DP			
	5856346	Jan 05, 2021	DS DP U-893			
	8658676	Oct 10, 2031	DP			
<u>CLINDAMYCIN PHOSPHATE - CLEOCIN</u>						
N 050767	001 6495157	Jul 20, 2020	DP			
<u>CLINDAMYCIN PHOSPHATE - CLINDAGEL</u>						
N 050782	001 6387383	Aug 03, 2020	DP U-818			
<u>CLINDAMYCIN PHOSPHATE - CLINDESSE</u>						
N 050793	001 6899890	Apr 27, 2023	DP U-137			
	9789057	Dec 02, 2026	DP U-137			
<u>CLINDAMYCIN PHOSPHATE - EVOCLIN</u>						
N 050801	001 7141237	Feb 03, 2024	DP			
	7374747	Jan 23, 2024	DP U-921			
<u>CLINDAMYCIN PHOSPHATE; TRETINOIN - ZIANA</u>						
N 050802	001 6387383	Aug 03, 2020	DP U-916			
<u>CLOBAZAM - SYMPAZAN</u>						
N 210833	001 8603514	Apr 03, 2024	DP			
	8765167	Feb 20, 2024	DP			
<u>CLOBAZAM - SYMPAZAN</u>						
N 210833	002 8603514	Apr 03, 2024	DP			
	8765167	Feb 20, 2024	DP			
<u>CLOBAZAM - SYMPAZAN</u>						
N 210833	003 8603514	Apr 03, 2024	DP			
	8765167	Feb 20, 2024	DP			

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<u>CLOBETASOL PROPIONATE - CLOBEX</u>						
N 021644	001 7700081	Jan 03, 2022	U-1044			
<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		NP	Nov 28, 2020
	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			
	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>CLOZAPINE - VERSACLOZ</u>						
N 203479	001 8057811	May 01, 2028	DP			
<u>COBICISTAT - TYBOST</u>						
N 203094	001 10039718	Oct 04, 2032	DP		NPP	Aug 22, 2022
	8148374	Sep 03, 2029	DS DP U-1279		ODE-260	Aug 22, 2026
<u>COBICISTAT; DARUNAVIR - PREZCOBIX</u>						
N 205395	001 10039718	Oct 04, 2032	DP			
	7700645	Dec 26, 2026	DS DP			
	7700645*PED	Jun 26, 2027				
	8148374	Sep 03, 2029	DS DP U-1279			
	8518987	Feb 16, 2024	DS DP			
	8518987*PED	Aug 16, 2024				
<u>COBICISTAT; DARUNAVIR; EMTRICITABINE; TENOFOVIR ALAFENAMIDE FUMARATE - SYMTUZA</u>						
N 210455	001 10039718	Oct 04, 2032	DP		NC	Jul 17, 2020
	6642245	Nov 04, 2020	U-2352		NCE	Nov 05, 2020
	6703396	Mar 09, 2021	DS DP			
	7390791	May 07, 2022	DS DP			
	7700645	Dec 26, 2026	DS DP			
	7803788	Feb 02, 2022	U-2352			
	8148374	Sep 03, 2029	DS DP U-2353			
	8148374	Sep 03, 2029	DS DP U-2364			
	8148374	Sep 03, 2029	DS DP U-2365			
	8518987	Feb 16, 2024	DS DP			
	8754065	Aug 15, 2032	DS DP U-2352			
	9296769	Aug 15, 2032	DS DP U-2352			
<u>COBICISTAT; ELVITEGRAVIR; EMTRICITABINE; TENOFOVIR ALAFENAMIDE FUMARATE - GENVOYA</u>						
N 207561	001 10039718	Oct 04, 2032	DP		D-173	Dec 10, 2021
	6642245	Nov 04, 2020	U-257		NCE	Nov 05, 2020
	6642245*PED	May 04, 2021			NPP	Sep 25, 2020
	6703396	Mar 09, 2021	DS DP			
	6703396*PED	Sep 09, 2021				
	7176220	Aug 27, 2026	DS DP U-257			
	7390791	May 07, 2022	DS DP			
	7635704	Oct 26, 2026	DS DP U-257			
	7803788	Feb 02, 2022	U-257			
	8148374	Sep 03, 2029	DS DP U-1279			
	8633219	Apr 24, 2030	DP U-257			
	8754065	Aug 15, 2032	DS DP U-257			
	8981103	Oct 26, 2026	DS DP			
	9296769	Aug 15, 2032	DS DP U-257			
	9891239	Sep 03, 2029	DP U-257			
<u>COBICISTAT; ELVITEGRAVIR; EMTRICITABINE; TENOFOVIR DISOPROXIL FUMARATE - STRIBILD</u>						
N 203100	001 10039718	Oct 04, 2032	DP		NPP	Jan 27, 2020
	6642245	Nov 04, 2020	U-257			
	6703396	Mar 09, 2021	DS DP			

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<u>COBICICISTAT; ELVITEGRAVIR; EMTRICITABINE; TENOFOVIR DISOPROXIL FUMARATE - STRIBILD</u>						
N 203100 001	7176220	Aug 27, 2026	DS DP U-257			
	7635704	Oct 26, 2026	DS DP U-257			
	8148374	Sep 03, 2029	DS DP U-1279			
	8592397	Jan 13, 2024	DP U-257			
	8633219	Apr 24, 2030	DP U-257			
	8716264	Jan 13, 2024	DP U-257			
	8981103	Oct 26, 2026	DS DP			
	9457036	Jan 13, 2024	DP U-257			
	9744181	Jan 13, 2024	DP U-257			
	9891239	Sep 03, 2029	DP U-257			
<u>COBIMETINIB FUMARATE - COTELLIC</u>						
N 206192 001	10478400	Jun 29, 2036	DS DP U-1776		NCE	Nov 10, 2020
	7803839	Feb 01, 2027	DS DP		ODE-101	Nov 10, 2022
	8362002	Oct 05, 2026	U-1776			
<u>COCAINE HYDROCHLORIDE - GOPRELTO</u>						
N 209963 001	10016407	Feb 07, 2037		U-2329	NCE	Dec 14, 2022
	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		
	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		
	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		
	9907751	Nov 22, 2036		DP		
<u>COLESEVELAM HYDROCHLORIDE - WELCHOL</u>						
N 021176 001	7229613	Apr 17, 2022		U-851		
<u>COLESEVELAM HYDROCHLORIDE - WELCHOL</u>						
N 022362 001	7229613	Apr 17, 2022		U-493		
<u>COLESEVELAM HYDROCHLORIDE - WELCHOL</u>						
N 022362 002	7229613	Apr 17, 2022		U-493		
<u>COLESEVELAM HYDROCHLORIDE - WELCHOL</u>						
N 210895 001	7229613	Apr 17, 2022		U-2516		

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<u>COPANLISIB DIHYDROCHLORIDE - ALIQOPA</u>						
N 209936 001	10383876	Mar 29, 2032	DS DP		NCE	Sep 14, 2022
	7511041	May 13, 2024	DS DP		ODE-155	Sep 14, 2024
	9636344	Mar 29, 2032		U-2124		
	RE46856	Oct 22, 2029	DS DP	U-2124		
<u>CRISABOROLE - EUCRISA</u>						
N 207695 001	8039451	Jun 11, 2026	DS DP		NCE	Dec 14, 2021
	8039451*PED	Dec 11, 2026			PED	Jun 14, 2022
	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		
	9682092*PED	Aug 16, 2027				
<u>CRIZOTINIB - XALKORI</u>						
N 202570 001	7230098	Aug 26, 2025	DS		ODE-111	Mar 11, 2023
	7825137	May 12, 2027		U-1179		
	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		ODE-111	Mar 11, 2023
	7825137	May 12, 2027		U-1179		
	7858643	Oct 08, 2029	DS DP			
	8217057	Nov 06, 2029	DS DP			
	8785632	Mar 01, 2025	DS			
<u>CROFELEMER - MYTESI</u>						
N 202292 001	7341744	Jun 02, 2022	DP	U-1319		
	8962680	Oct 31, 2031		U-1319		
	9585868	Oct 31, 2031		U-1319		
<u>CYANOCOBALAMIN - NASCOBAL</u>						
N 021642 001	7229636	Aug 01, 2024	DP	U-817		
	7404489	Mar 12, 2024	DP			
	7879349	Aug 01, 2024	DP	U-1152		
	8003353	Aug 01, 2024		U-817		
	8940714	Feb 26, 2024		U-1152		
	9415007	Jul 28, 2024		U-1896		
<u>CYCLOBENZAPRINE HYDROCHLORIDE - AMRIX</u>						
N 021777 001	7387793	Feb 26, 2025	DP			
	7544372	Nov 14, 2023		U-979		
	7790199	Nov 14, 2023	DP			
	7820203	Nov 14, 2023	DP			
	7829121	Nov 14, 2023		U-1088		
	8877245	Nov 14, 2023		U-979		
	9375410	Nov 14, 2023		U-1088		
	9399025	Nov 14, 2023	DP	U-979		
<u>CYCLOBENZAPRINE HYDROCHLORIDE - AMRIX</u>						
N 021777 002	7387793	Feb 26, 2025	DP			
	7544372	Nov 14, 2023		U-979		
	7790199	Nov 14, 2023	DP			
	7820203	Nov 14, 2023	DP			
	7829121	Nov 14, 2023		U-1088		
	8877245	Nov 14, 2023		U-979		
	9375410	Nov 14, 2023		U-1088		
	9399025	Nov 14, 2023	DP	U-979		
<u>CYCLOSPORINE - RESTASIS</u>						
N 050790 001	8629111	Aug 27, 2024	DP			
	8633162	Aug 27, 2024		U-1479		
	8642556	Aug 27, 2024	DP			
	8648048	Aug 27, 2024		U-1483		
	8685930	Aug 27, 2024	DP			
	9248191	Aug 27, 2024		U-1479		

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<u>CYCLOSPORINE - RESTASIS MULTIDOSE</u>						
N 050790 002	8292129	Feb 25, 2031	DP			
	8561859	Apr 16, 2032	DP			
	8629111	Aug 27, 2024	DP			
	8633162	Aug 27, 2024		U-1479		
	8642556	Aug 27, 2024	DP			
	8648048	Aug 27, 2024		U-1483		
	8685930	Aug 27, 2024	DP			
	9248191	Aug 27, 2024		U-1479		
	9669974	May 11, 2034	DP			
	9676525	Feb 07, 2034	DP			
<u>CYCLOSPORINE - CEQUA</u>						
N 210913 001	10441630	Aug 23, 2033	DP			
	8980839	Aug 23, 2033	DP	U-1483		
	9937225	Aug 23, 2033	DP	U-1483		
<u>CYSTEAMINE BITARTRATE - PROCYSBI</u>						
N 203389 001	10143665	Aug 16, 2036		U-1399	M-216	Dec 22, 2020
	10328037	Aug 16, 2036		U-1399	ODE-162	Dec 22, 2024
	10548859	Aug 16, 2036		U-1399	ODE-45	Apr 30, 2020
	8026284	Sep 22, 2027		U-1399	ODE-97	Aug 14, 2022
	8026284*PED	Mar 22, 2028			PED	Oct 30, 2020
	9173851	Jun 17, 2034	DP		PED	Jun 22, 2021
	9173851*PED	Dec 17, 2034			PED	Feb 14, 2023
	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		
	9925157	Jan 26, 2027	DS DP	U-1399		
	9925158	Jan 26, 2027	DS DP	U-1399		
	<u>CYSTEAMINE BITARTRATE - PROCYSBI</u>					
N 203389 002	10143665	Aug 16, 2036		U-1399	M-216	Dec 22, 2020
	10328037	Aug 16, 2036		U-1399	ODE-162	Dec 22, 2024
	10548859	Aug 16, 2036		U-1399	ODE-45	Apr 30, 2020
	8026284	Sep 22, 2027		U-1399	ODE-97	Aug 14, 2022
	8026284*PED	Mar 22, 2028			PED	Oct 30, 2020
	9173851	Jun 17, 2034	DP		PED	Jun 22, 2021
	9173851*PED	Dec 17, 2034			PED	Feb 14, 2023
	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		
	9925157	Jan 26, 2027	DS DP	U-1399		
	9925158	Jan 26, 2027	DS DP	U-1399		
	<u>CYSTEAMINE BITARTRATE - PROCYSBI</u>					
N 213491 001	10143665	Aug 16, 2036		U-1399		
	10328037	Aug 16, 2036		U-1399		
	10548859	Aug 16, 2036		U-1399		
	8026284	Sep 22, 2027		U-1399		
	9173851	Jun 17, 2034	DP			
	9192590	Jan 26, 2027		U-1399		
	9198882	Jan 26, 2027		U-1399		
	9233077	Jun 17, 2034	DP			
	9925156	Jan 26, 2027	DP	U-1399		
	9925157	Jan 26, 2027	DP	U-1399		
9925158	Jan 26, 2027	DP	U-1399			
<u>CYSTEAMINE BITARTRATE - PROCYSBI</u>						
N 213491 002	10143665	Aug 16, 2036		U-1399		
	10328037	Aug 16, 2036		U-1399		
	10548859	Aug 16, 2036		U-1399		
	8026284	Sep 22, 2027		U-1399		

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<u>CYSTEAMINE BITARTRATE - PROCYSBI</u>						
N 213491	002	9173851	Jun 17, 2034	DP		
		9192590	Jan 26, 2027	U-1399		
		9198882	Jan 26, 2027	U-1399		
		9233077	Jun 17, 2034	DP		
		9925156	Jan 26, 2027	DP U-1399		
		9925157	Jan 26, 2027	DP U-1399		
		9925158	Jan 26, 2027	DP U-1399		
<u>CYSTEINE HYDROCHLORIDE - ELCYS</u>						
N 210660	001	10478453	Jan 15, 2039	DP U-2752		
		10583155	Jan 15, 2039	DP U-2752		
<u>CYSTEINE HYDROCHLORIDE - NOURESS</u>						
N 212535	001	10493051	Mar 15, 2039	DP		
		10543186	Mar 15, 2039	U-2722		
<u>CYTARABINE; DAUNORUBICIN - VYXEOS</u>						
N 209401	001	10028912	Sep 29, 2034	DP U-2341	NP	Aug 03, 2020
		10028912	Sep 29, 2034	DP U-2342		
		10166184	Oct 15, 2032	DP U-2341		
		7850990	Jan 23, 2027	DP U-2090		
		8022279	Sep 14, 2027	DP U-2090		
		8092828	Apr 01, 2029	U-2090		
		8431806	Apr 22, 2025	DP U-2090		
		8518437	Jun 07, 2026	DP		
		9271931	Jan 23, 2027	DP		
<u>DABIGATRAN ETEXILATE MESYLATE - PRADAXA</u>						
N 022512	001	6087380	Dec 28, 2021	DS DP U-1931		
		7866474	Aug 31, 2027	DP	Y	
		7932273	Sep 07, 2025	DS DP		
		9034822	Jan 20, 2031	U-1759		
		9925174	Jun 14, 2023	DP		
<u>DABIGATRAN ETEXILATE MESYLATE - PRADAXA</u>						
N 022512	002	6087380	Dec 28, 2021	DS DP U-1931		
		7866474	Aug 31, 2027	DP	Y	
		7932273	Sep 07, 2025	DS DP		
		9034822	Jan 20, 2031	U-1759		
		9925174	Jun 14, 2023	DP		
<u>DABIGATRAN ETEXILATE MESYLATE - PRADAXA</u>						
N 022512	003	6087380	Dec 28, 2021	DS DP U-1931		
		7866474	Aug 31, 2027	DP	Y	
		7932273	Sep 07, 2025	DS DP		
		9034822	Jan 20, 2031	U-1759		
		9925174	Jun 14, 2023	DP		
<u>DABRAFENIB MESYLATE - TAFINLAR</u>						
N 202806	001	7994185	Jan 20, 2030	DS DP U-1406	I-745	Jun 22, 2020
		7994185	Jan 20, 2030	DS DP U-2031	I-778	Apr 30, 2021
		7994185	Jan 20, 2030	DS DP U-2032	I-781	May 04, 2021
		7994185	Jan 20, 2030	DS DP U-2296	M-246	Oct 06, 2022
		8415345	Jan 20, 2030	DS DP U-1406	ODE-147	Jun 22, 2024
		8415345	Jan 20, 2030	DS DP U-2031	ODE-182	Apr 30, 2025
		8415345	Jan 20, 2030	DS DP U-2032	ODE-183	May 04, 2025
		8415345	Jan 20, 2030	DS DP U-2296	ODE-47	May 29, 2020
		8703781	Oct 15, 2030	DS DP U-1713	ODE-58	Jan 09, 2021
		8703781	Oct 15, 2030	DS DP U-2032		
		8703781	Oct 15, 2030	DS DP U-2296		
		8703781	Oct 15, 2030	DS DP U-2297		
		8703781	Oct 15, 2030	DS DP U-2298		
		8835443	Jun 10, 2025	U-2026		
		8835443	Jun 10, 2025	U-2027		
		8835443	Jun 10, 2025	U-2296		
		8835443	Jun 10, 2025	U-2298		
		8952018	Oct 15, 2030	U-2027		
		9233956	May 04, 2029	U-1811		
		9233956	May 04, 2029	U-2031		
		9233956	May 04, 2029	U-2032		

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<u>DABRAFENIB MESYLATE - TAFINLAR</u>						
N 202806 001	9233956	May 04, 2029	U-2296			
	9233956	May 04, 2029	U-2297			
<u>DABRAFENIB MESYLATE - TAFINLAR</u>						
N 202806 002	7994185	Jan 20, 2030	DS DP U-1406		I-745	Jun 22, 2020
	7994185	Jan 20, 2030	DS DP U-2031		I-778	Apr 30, 2021
	7994185	Jan 20, 2030	DS DP U-2032		I-781	May 04, 2021
	7994185	Jan 20, 2030	DS DP U-2296		M-246	Oct 06, 2022
	8415345	Jan 20, 2030	DS DP U-1406		ODE-147	Jun 22, 2024
	8415345	Jan 20, 2030	DS DP U-2031		ODE-182	Apr 30, 2025
	8415345	Jan 20, 2030	DS DP U-2032		ODE-183	May 04, 2025
	8415345	Jan 20, 2030	DS DP U-2296		ODE-47	May 29, 2020
	8703781	Oct 15, 2030	DS DP U-1713		ODE-58	Jan 09, 2021
	8703781	Oct 15, 2030	DS DP U-2032			
	8703781	Oct 15, 2030	DS DP U-2296			
	8703781	Oct 15, 2030	DS DP U-2297			
	8703781	Oct 15, 2030	DS DP U-2298			
	8835443	Jun 10, 2025	U-2026			
	8835443	Jun 10, 2025	U-2027			
	8835443	Jun 10, 2025	U-2296			
	8835443	Jun 10, 2025	U-2298			
	8952018	Oct 15, 2030	U-2027			
	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	May 04, 2029	U-2297			
<u>DACLATASVIR DIHYDROCHLORIDE - DAKLINZA</u>						
N 206843 001	8329159	Jul 24, 2029	DS		NCE	Jul 24, 2020
	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		NCE	Jul 24, 2020
	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	7772243	Aug 26, 2028	DS DP		NCE	Sep 27, 2023
	8623883	May 05, 2025	U-1403		ODE-206	Sep 27, 2025
					ODE-213	Sep 27, 2025
<u>DACOMITINIB - VIZIMPRO</u>						
N 211288 002	7772243	Aug 26, 2028	DS DP		NCE	Sep 27, 2023
	8623883	May 05, 2025	U-1403		ODE-206	Sep 27, 2025
					ODE-213	Sep 27, 2025
<u>DACOMITINIB - VIZIMPRO</u>						
N 211288 003	7772243	Aug 26, 2028	DS DP		NCE	Sep 27, 2023
	8623883	May 05, 2025	U-1403		ODE-206	Sep 27, 2025
					ODE-213	Sep 27, 2025
<u>DALBAVANCIN HYDROCHLORIDE - DALVANCE</u>						
N 021883 001	6900175	Dec 25, 2023	U-1517		NCE	May 23, 2019
	7115564	Nov 14, 2023	DP		GAIN	May 23, 2024

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APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>DALBAVANCIN HYDROCHLORIDE - DALVANCE</u>						
N 021883 001	7119061	Nov 14, 2023	DP			
	8143212	Nov 14, 2023	U-1517			
<u>DALFAMPRIDINE - AMPYRA</u>						
N 022250 001	8007826	May 26, 2027	U-1030	Y		
	8354437	Dec 22, 2026	U-1030	Y		
	8440703	Apr 08, 2025	U-1030	Y		
<u>DALTEPARIN SODIUM - FRAGMIN</u>						
N 020287 001					NPP	May 16, 2022
<u>DALTEPARIN SODIUM - FRAGMIN</u>						
N 020287 002					NPP	May 16, 2022
<u>DALTEPARIN SODIUM - FRAGMIN</u>						
N 020287 003					NPP	May 16, 2022
<u>DALTEPARIN SODIUM - FRAGMIN</u>						
N 020287 004					NPP	May 16, 2022
<u>DALTEPARIN SODIUM - FRAGMIN</u>						
N 020287 005					NPP	May 16, 2022
<u>DALTEPARIN SODIUM - FRAGMIN</u>						
N 020287 006					NPP	May 16, 2022
<u>DALTEPARIN SODIUM - FRAGMIN</u>						
N 020287 007					NPP	May 16, 2022
<u>DALTEPARIN SODIUM - FRAGMIN</u>						
N 020287 008					NPP	May 16, 2022
<u>DALTEPARIN SODIUM - FRAGMIN</u>						
N 020287 009					NPP	May 16, 2022
<u>DALTEPARIN SODIUM - FRAGMIN</u>						
N 020287 010					NPP	May 16, 2022
<u>DALTEPARIN SODIUM - FRAGMIN</u>						
N 020287 011					NPP	May 16, 2022
<u>DANTROLENE SODIUM - RYANODEX</u>						
N 205579 001	7758890	Jul 01, 2025	DP		ODE-69	Jul 22, 2021
	8110225	Dec 24, 2022	DP			
	8604072	Dec 24, 2022	DP			
	8685460	Feb 15, 2023	U-1546			
	9884044	Jun 13, 2022	DP U-1546			
<u>DAPAGLIFLOZIN - FARXIGA</u>						
N 202293 001	6414126	Oct 04, 2020	DS DP U-2139		M-212	Oct 20, 2020
	6414126	Oct 04, 2020	DS DP U-493		M-238	Feb 22, 2022
	6515117	Oct 04, 2020	DS DP U-2139			
	6515117	Oct 04, 2020	DS DP U-493			
	6936590	Oct 04, 2020	U-493			
	7456254	Jun 30, 2025	DP U-2139			
	7851502	Aug 19, 2028	DP			
	7919598	Dec 16, 2029	DS			
	8221786	Mar 21, 2028	DP			
	8329648	Aug 18, 2026	U-2139			
	8329648	Aug 18, 2026	U-2212			
	8329648	Aug 18, 2026	U-2213			
	8361972	Mar 21, 2028	U-2139			
	8361972	Mar 21, 2028	U-493			
	8431685	Apr 13, 2025	DP U-2139			
	8461105	Apr 13, 2025	DP U-2139			
	8501698	Jun 20, 2027	DP U-493			
	8685934	May 26, 2030	U-1522			
	8716251	Mar 21, 2028	DP			
	8721615	Jan 18, 2030	DP	Y		
	8906851	Aug 18, 2026	U-2139			
	9198925	Oct 04, 2020	U-2139			

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<u>DAPAGLIFLOZIN - FARXIGA</u>						
N 202293 001	9198925	Oct 04, 2020	U-493			
	9238076	Apr 15, 2024	DP U-2139			
<u>DAPAGLIFLOZIN - FARXIGA</u>						
N 202293 002	6414126	Oct 04, 2020	DS DP U-2139		M-212	Oct 20, 2020
	6414126	Oct 04, 2020	DS DP U-493		M-238	Feb 22, 2022
	6515117	Oct 04, 2020	DS DP U-2139			
	6515117	Oct 04, 2020	DS DP U-493			
	6936590	Oct 04, 2020	U-493			
	7456254	Jun 30, 2025	DP U-2139			
	7851502	Aug 19, 2028	DP			
	7919598	Dec 16, 2029	DS			
	8221786	Mar 21, 2028	DP			
	8329648	Aug 18, 2026	U-2139			
	8329648	Aug 18, 2026	U-2212			
	8329648	Aug 18, 2026	U-2213			
	8361972	Mar 21, 2028	U-2139			
	8361972	Mar 21, 2028	U-493			
	8431685	Apr 13, 2025	DP U-2139			
	8461105	Apr 13, 2025	DP U-2139			
	8501698	Jun 20, 2027	DP U-493			
	8685934	May 26, 2030	U-1522			
	8716251	Mar 21, 2028	DP			
	8721615	Jan 18, 2030	DP	Y		
	8906851	Aug 18, 2026	U-2139			
	9198925	Oct 04, 2020	U-2139			
	9198925	Oct 04, 2020	U-493			
	9238076	Apr 15, 2024	DP U-2139			
<u>DAPAGLIFLOZIN; METFORMIN HYDROCHLORIDE - XIGDUO XR</u>						
N 205649 001	6414126	Oct 04, 2020	DS DP U-493			
	6515117	Oct 04, 2020	DS DP U-493			
	6936590	Oct 04, 2020	U-493			
	7919598	Dec 16, 2029	DS			
	8501698	Jun 20, 2027	DP U-493			
	8685934	May 26, 2030	U-1522			
	9198925	Oct 04, 2020	U-493			
	9616028	Nov 12, 2030	DP			
<u>DAPAGLIFLOZIN; METFORMIN HYDROCHLORIDE - XIGDUO XR</u>						
N 205649 002	6414126	Oct 04, 2020	DS DP U-493			
	6515117	Oct 04, 2020	DS DP U-493			
	6936590	Oct 04, 2020	U-493			
	7919598	Dec 16, 2029	DS			
	8501698	Jun 20, 2027	DP U-493			
	8685934	May 26, 2030	U-1522			
	9198925	Oct 04, 2020	U-493			
	9616028	Nov 12, 2030	DP			
<u>DAPAGLIFLOZIN; METFORMIN HYDROCHLORIDE - XIGDUO XR</u>						
N 205649 003	6414126	Oct 04, 2020	DS DP U-493			
	6515117	Oct 04, 2020	DS DP U-493			
	6936590	Oct 04, 2020	U-493			
	7919598	Dec 16, 2029	DS			
	8501698	Jun 20, 2027	DP U-493			
	8685934	May 26, 2030	U-1522			
	9198925	Oct 04, 2020	U-493			
	9616028	Nov 12, 2030	DP			
<u>DAPAGLIFLOZIN; METFORMIN HYDROCHLORIDE - XIGDUO XR</u>						
N 205649 004	6414126	Oct 04, 2020	DS DP U-493			
	6515117	Oct 04, 2020	DS DP U-493			
	6936590	Oct 04, 2020	U-493			
	7919598	Dec 16, 2029	DS			
	8501698	Jun 20, 2027	DP U-493			
	8685934	May 26, 2030	U-1522			
	9198925	Oct 04, 2020	U-493			
	9616028	Nov 12, 2030	DP			

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<u>DAPAGLIFLOZIN; METFORMIN HYDROCHLORIDE - XIGDUO XR</u>						
N 205649 005	6414126	Oct 04, 2020	DS DP U-493			
	6515117	Oct 04, 2020	DS DP U-493			
	6936590	Oct 04, 2020	U-493			
	7919598	Dec 16, 2029	DS			
	8501698	Jun 20, 2027	DP U-493			
	8685934	May 26, 2030	U-1522			
	9198925	Oct 04, 2020	U-493			
	9616028	Nov 12, 2030	DP			
<u>DAPAGLIFLOZIN; METFORMIN HYDROCHLORIDE; SAXAGLIPTIN HYDROCHLORIDE - QTERNMET XR</u>						
N 210874 001	6414126	Oct 04, 2020	DS DP U-493		NP	May 02, 2022
	6515117	Oct 04, 2020	DS DP U-493			
	6936590	Oct 04, 2020	U-493			
	7919598	Dec 16, 2029	DS			
	8501698	Jun 20, 2027	DP U-493			
	8628799	Jul 13, 2025	DP			
	8716251	Mar 21, 2028	DP			
	9198925	Oct 04, 2020	U-493			
	9616028	Nov 12, 2030	DP			
	RE44186	Jul 31, 2023	DS DP U-493			
<u>DAPAGLIFLOZIN; METFORMIN HYDROCHLORIDE; SAXAGLIPTIN HYDROCHLORIDE - QTERNMET XR</u>						
N 210874 002	6414126	Oct 04, 2020	DS DP U-493		NP	May 02, 2022
	6515117	Oct 04, 2020	DS DP U-493			
	6936590	Oct 04, 2020	U-493			
	7919598	Dec 16, 2029	DS			
	8501698	Jun 20, 2027	DP U-493			
	8628799	Jul 13, 2025	DP			
	8716251	Mar 21, 2028	DP			
	9198925	Oct 04, 2020	U-493			
	9616028	Nov 12, 2030	DP			
	RE44186	Jul 31, 2023	DS DP U-493			
<u>DAPAGLIFLOZIN; METFORMIN HYDROCHLORIDE; SAXAGLIPTIN HYDROCHLORIDE - QTERNMET XR</u>						
N 210874 003	6414126	Oct 04, 2020	DS DP U-493		NP	May 02, 2022
	6515117	Oct 04, 2020	DS DP U-493			
	6936590	Oct 04, 2020	U-493			
	7919598	Dec 16, 2029	DS			
	8501698	Jun 20, 2027	DP U-493			
	8628799	Jul 13, 2025	DP			
	8716251	Mar 21, 2028	DP			
	9198925	Oct 04, 2020	U-493			
	9616028	Nov 12, 2030	DP			
	RE44186	Jul 31, 2023	DS DP U-493			
<u>DAPAGLIFLOZIN; METFORMIN HYDROCHLORIDE; SAXAGLIPTIN HYDROCHLORIDE - QTERNMET XR</u>						
N 210874 004	6414126	Oct 04, 2020	DS DP U-493		NP	May 02, 2022
	6515117	Oct 04, 2020	DS DP U-493			
	6936590	Oct 04, 2020	U-493			
	7919598	Dec 16, 2029	DS			
	8501698	Jun 20, 2027	DP U-493			
	8628799	Jul 13, 2025	DP			
	8716251	Mar 21, 2028	DP			
	9198925	Oct 04, 2020	U-493			
	9616028	Nov 12, 2030	DP			
	RE44186	Jul 31, 2023	DS DP U-493			
<u>DAPAGLIFLOZIN; SAXAGLIPTIN HYDROCHLORIDE - QTERN</u>						
N 209091 001	6414126	Oct 04, 2020	DS DP U-493		I-804	May 02, 2022
	6515117	Oct 04, 2020	DS DP U-493		NC	Feb 27, 2020
	6936590	Oct 04, 2020	U-1976			
	6936590	Oct 04, 2020	U-1977			
	6936590	Oct 04, 2020	U-493			
	7919598	Dec 16, 2029	DS			
	8221786	Mar 21, 2028	DP			
	8361972	Mar 21, 2028	U-1976			
	8361972	Mar 21, 2028	U-1977			
	8361972	Mar 21, 2028	U-493			
	8501698	Jun 20, 2027	DP U-1976			
	8501698	Jun 20, 2027	DP U-1977			

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<u>DAPAGLIFLOZIN; SAXAGLIPTIN HYDROCHLORIDE - QTERN</u>						
N 209091 001	8501698	Jun 20, 2027	DP U-493			
	8628799	Jul 13, 2025	DP			
	8716251	Mar 21, 2028	DP			
	9198925	Oct 04, 2020	U-1976			
	9198925	Oct 04, 2020	U-1977			
	9198925	Oct 04, 2020	U-493			
	RE44186	Jul 31, 2023	DS DP U-493			
<u>DAPAGLIFLOZIN; SAXAGLIPTIN HYDROCHLORIDE - QTERN</u>						
N 209091 002	6414126	Oct 04, 2020	DS DP U-493		NS	May 02, 2022
	6515117	Oct 04, 2020	DS DP U-493			
	6936590	Oct 04, 2020	U-493			
	7919598	Dec 16, 2029	DS			
	8221786	Mar 21, 2028	DP			
	8361972	Mar 21, 2028	U-493			
	8501698	Jun 20, 2027	DP U-493			
	8628799	Jul 13, 2025	DP			
	8716251	Mar 21, 2028	DP			
	9198925	Oct 04, 2020	U-493			
	RE44186	Jul 31, 2023	DS DP U-493			
<u>DAPSONE - ACZONE</u>						
N 207154 001	9161926	Nov 18, 2033	DP		NPP	Sep 10, 2022
	9517219	Nov 18, 2033	U-1033			
<u>DAPTOMYCIN - DAPTOMYCIN</u>						
A 212667 001					CGT	Jan 12, 2020
<u>DAPTOMYCIN - CUBICIN</u>						
N 021572 002	8003673	Sep 04, 2028	U-1180		M-211 NPP	Sep 01, 2020 Mar 29, 2020
<u>DAPTOMYCIN - CUBICIN RF</u>						
N 021572 003	9138456	Nov 23, 2030	DP		M-211 NPP	Sep 01, 2020 Mar 29, 2020
<u>DAROLUTAMIDE - NUBEQA</u>						
N 212099 001	10010530	Jan 28, 2036	DS		NCE	Jul 30, 2024
	10383853	Jan 28, 2036	DS			
	8975254	Oct 27, 2030	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			
	8518987	Feb 16, 2024	DS DP			
	8518987*PED	Aug 16, 2024				
<u>DARUNAVIR - PREZISTA</u>						
N 021976 002	7700645	Dec 26, 2026	DS DP			
	8518987	Feb 16, 2024	DS DP			
	8518987*PED	Aug 16, 2024				
<u>DARUNAVIR - PREZISTA</u>						
N 021976 003	7700645	Dec 26, 2026	DS DP			
	8518987	Feb 16, 2024	DS DP			
	8518987*PED	Aug 16, 2024				
<u>DARUNAVIR - PREZISTA</u>						
N 021976 004	7700645	Dec 26, 2026	DS DP			
	8518987	Feb 16, 2024	DS DP			
	8518987*PED	Aug 16, 2024				
<u>DARUNAVIR - PREZISTA</u>						
N 021976 005	7700645	Dec 26, 2026	DS DP			
	8518987	Feb 16, 2024	DS DP			
	8518987*PED	Aug 16, 2024				
<u>DARUNAVIR - PREZISTA</u>						
N 021976 006	7700645	Dec 26, 2026	DS DP			
	8518987	Feb 16, 2024	DS DP			

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<u>DARUNAVIR - PREZISTA</u>						
N 021976	006 8518987*PED	Aug 16, 2024				
<u>DARUNAVIR - PREZISTA</u>						
N 202895	001 7700645	Dec 26, 2026	DS DP			
	8518987	Feb 16, 2024	DS DP			
	8518987*PED	Aug 16, 2024				
<u>DASABUVIR SODIUM; OMBITASVIR; PARITAPREVIR; RITONAVIR - VIEKIRA PAK (COPACKAGED)</u>						
N 206619	001 10201542	Oct 18, 2033	DP U-1753			
	7364752	Nov 10, 2020	DP			
	8188104	May 17, 2029	DS DP U-1636			
	8268349	Aug 25, 2024	DP			
	8399015	Aug 25, 2024	DP			
	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			
	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			
	7364752	Nov 10, 2020	DP			
	8188104	May 17, 2029	DS DP U-1636			
	8268349	Aug 25, 2024	DP			
	8399015	Aug 25, 2024	DP			
	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 - SPRYCEL</u>						
N 021986	001 6596746	Jun 28, 2020	DS DP U-748		I-791	Dec 21, 2021
	6596746	Jun 28, 2020	DS DP U-780		NPP	Nov 09, 2020
	6596746*PED	Dec 28, 2020			ODE-164	Nov 09, 2024
	7125875	Apr 13, 2020	U-779		ODE-225	Dec 21, 2025
	7125875	Apr 13, 2020	U-780		PED	May 09, 2021
	7125875*PED	Oct 13, 2020			PED	Jun 21, 2022
	7153856	Apr 28, 2020	U-780		PED	May 09, 2025
	7153856*PED	Oct 28, 2020			PED	Jun 21, 2026
	7491725	Mar 28, 2026	DS DP			
	7491725*PED	Sep 28, 2026				
	8680103	Feb 04, 2025	DP			
	8680103*PED	Aug 04, 2025				
<u>DASATINIB - SPRYCEL</u>						
N 021986	002 6596746	Jun 28, 2020	DS DP U-748		I-791	Dec 21, 2021
	6596746	Jun 28, 2020	DS DP U-780		NPP	Nov 09, 2020
	6596746*PED	Dec 28, 2020			ODE-164	Nov 09, 2024
	7125875	Apr 13, 2020	U-779		ODE-225	Dec 21, 2025
	7125875	Apr 13, 2020	U-780		PED	May 09, 2021

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<u>DASATINIB - SPRYCEL</u>						
N 021986 002	7125875*PED	Oct 13, 2020			PED	Jun 21, 2022
	7153856	Apr 28, 2020	U-780		PED	May 09, 2025
	7153856*PED	Oct 28, 2020			PED	Jun 21, 2026
	7491725	Mar 28, 2026	DS DP			
	7491725*PED	Sep 28, 2026				
	8680103	Feb 04, 2025	DP			
	8680103*PED	Aug 04, 2025				
<u>DASATINIB - SPRYCEL</u>						
N 021986 003	6596746	Jun 28, 2020	DS DP U-748		I-791	Dec 21, 2021
	6596746	Jun 28, 2020	DS DP U-780		NPP	Nov 09, 2020
	6596746*PED	Dec 28, 2020			ODE-164	Nov 09, 2024
	7125875	Apr 13, 2020	U-779		ODE-225	Dec 21, 2025
	7125875	Apr 13, 2020	U-780		PED	May 09, 2021
	7125875*PED	Oct 13, 2020			PED	Jun 21, 2022
	7153856	Apr 28, 2020	U-780		PED	May 09, 2025
	7153856*PED	Oct 28, 2020			PED	Jun 21, 2026
	7491725	Mar 28, 2026	DS DP			
	7491725*PED	Sep 28, 2026				
	8680103	Feb 04, 2025	DP			
	8680103*PED	Aug 04, 2025				
	<u>DASATINIB - SPRYCEL</u>					
N 021986 004	6596746	Jun 28, 2020	DS DP U-748		I-791	Dec 21, 2021
	6596746	Jun 28, 2020	DS DP U-780		NPP	Nov 09, 2020
	6596746*PED	Dec 28, 2020			ODE-164	Nov 09, 2024
	7125875	Apr 13, 2020	U-779		ODE-225	Dec 21, 2025
	7125875	Apr 13, 2020	U-780		PED	May 09, 2021
	7125875*PED	Oct 13, 2020			PED	Jun 21, 2022
	7153856	Apr 28, 2020	U-780		PED	May 09, 2025
	7153856*PED	Oct 28, 2020			PED	Jun 21, 2026
	7491725	Mar 28, 2026	DS DP			
	7491725*PED	Sep 28, 2026				
	8680103	Feb 04, 2025	DP			
	8680103*PED	Aug 04, 2025				
	<u>DASATINIB - SPRYCEL</u>					
N 021986 005	6596746	Jun 28, 2020	DS DP U-748		I-791	Dec 21, 2021
	6596746	Jun 28, 2020	DS DP U-780		NPP	Nov 09, 2020
	6596746*PED	Dec 28, 2020			ODE-164	Nov 09, 2024
	7125875	Apr 13, 2020	U-779		ODE-225	Dec 21, 2025
	7125875	Apr 13, 2020	U-780		PED	May 09, 2021
	7125875*PED	Oct 13, 2020			PED	Jun 21, 2022
	7153856	Apr 28, 2020	U-780		PED	May 09, 2025
	7153856*PED	Oct 28, 2020			PED	Jun 21, 2026
	7491725	Mar 28, 2026	DS DP			
	7491725*PED	Sep 28, 2026				
	8680103	Feb 04, 2025	DP			
	8680103*PED	Aug 04, 2025				
	<u>DASATINIB - SPRYCEL</u>					
N 021986 006	6596746	Jun 28, 2020	DS DP U-748		I-791	Dec 21, 2021
	6596746	Jun 28, 2020	DS DP U-780		NPP	Nov 09, 2020
	6596746*PED	Dec 28, 2020			ODE-164	Nov 09, 2024
	7125875	Apr 13, 2020	U-779		ODE-225	Dec 21, 2025
	7125875	Apr 13, 2020	U-780		PED	May 09, 2021
	7125875*PED	Oct 13, 2020			PED	Jun 21, 2022
	7153856	Apr 28, 2020	U-780		PED	May 09, 2025
	7153856*PED	Oct 28, 2020			PED	Jun 21, 2026
	7491725	Mar 28, 2026	DS DP			
	7491725*PED	Sep 28, 2026				
	8680103	Feb 04, 2025	DP			
	8680103*PED	Aug 04, 2025				
	<u>DEFERASIROX - DEFERASIROX</u>					
A 208697 002					PC	Jun 14, 2020

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<u>DEFERASIROX - EXJADE</u>						
N 021882	001				M-239 M-241 ODE-39	Dec 12, 2021 Jul 24, 2022 Jan 23, 2020
<u>DEFERASIROX - EXJADE</u>						
N 021882	002				M-239 M-241 ODE-39	Dec 12, 2021 Jul 24, 2022 Jan 23, 2020
<u>DEFERASIROX - EXJADE</u>						
N 021882	003				M-239 M-241 ODE-39	Dec 12, 2021 Jul 24, 2022 Jan 23, 2020
<u>DEFERASIROX - JADENU</u>						
N 206910	001 9283209	Nov 21, 2034	DS DP		M-239 M-241 ODE-39	Dec 12, 2021 Jul 24, 2022 Jan 23, 2020
<u>DEFERASIROX - JADENU</u>						
N 206910	002 9283209	Nov 21, 2034	DS DP		M-239 M-241 ODE-39	Dec 12, 2021 Jul 24, 2022 Jan 23, 2020
<u>DEFERASIROX - JADENU</u>						
N 206910	003 9283209	Nov 21, 2034	DS DP		M-239 M-241 ODE-39	Dec 12, 2021 Jul 24, 2022 Jan 23, 2020
<u>DEFERASIROX - JADENU SPRINKLE</u>						
N 207968	001				M-239 M-241	Dec 12, 2021 Jul 24, 2022
<u>DEFERASIROX - JADENU SPRINKLE</u>						
N 207968	002				M-239 M-241	Dec 12, 2021 Jul 24, 2022
<u>DEFERASIROX - JADENU SPRINKLE</u>						
N 207968	003				M-239 M-241	Dec 12, 2021 Jul 24, 2022
<u>DEFERIPRONE - FERRIPROX</u>						
N 021825	001 7049328	Jun 28, 2021		U-735		
<u>DEFERIPRONE - FERRIPROX</u>						
N 021825	002 7049328	Jun 28, 2021		U-735		
<u>DEFERIPRONE - FERRIPROX</u>						
N 208030	001 7049328 8703156	Jun 28, 2021 Oct 29, 2029		U-735 DP U-735		
<u>DEFERIPRONE - FERRIPROX</u>						
N 208030	002 7049328 8703156	Jun 28, 2021 Oct 29, 2029		U-735 DP U-735		
<u>DEFIBROTIDE SODIUM - DEFITELIO</u>						
N 208114	001				NCE ODE-112	Mar 30, 2021 Mar 30, 2023
<u>DEFLAZACORT - EMFLAZA</u>						
N 208684	001				NCE ODE-130 ODE-252	Feb 09, 2022 Feb 09, 2024 Jun 07, 2026
<u>DEFLAZACORT - EMFLAZA</u>						
N 208684	002				NCE ODE-130 ODE-252	Feb 09, 2022 Feb 09, 2024 Jun 07, 2026

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<u>DEFLAZACORT - EMFLAZA</u>						
N 208684	003				NCE ODE-130 ODE-252	Feb 09, 2022 Feb 09, 2024 Jun 07, 2026
<u>DEFLAZACORT - EMFLAZA</u>						
N 208684	004				NCE ODE-130 ODE-252	Feb 09, 2022 Feb 09, 2024 Jun 07, 2026
<u>DEFLAZACORT - EMFLAZA</u>						
N 208685	001				NCE ODE-130 ODE-252	Feb 09, 2022 Feb 09, 2024 Jun 07, 2026
<u>DEGARELIX ACETATE - FIRMAGON</u>						
N 022201	001	5925730	May 18, 2021	DS DP U-943		
		9415085	Apr 27, 2032	U-1895		
		9579359	Feb 10, 2029	U-1978		
<u>DEGARELIX ACETATE - FIRMAGON</u>						
N 022201	002	5925730	May 18, 2021	DS DP U-943		
		9415085	Apr 27, 2032	U-1895		
		9579359	Feb 10, 2029	U-1978		
<u>DELAFLORACIN MEGLUMINE - BAXDELA</u>						
N 208610	001	7728143	Nov 20, 2027	DS	I-815	Oct 24, 2022
		8252813	Oct 02, 2026	DP U-2028	NCE	Jun 19, 2022
		8273892	Aug 06, 2026	DS	GAIN	Jun 19, 2027
		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>DELAFLORACIN MEGLUMINE - BAXDELA</u>						
N 208611	001	7635773	Mar 13, 2029	DP	I-815	Oct 24, 2022
		7728143	Nov 20, 2027	DS	NCE	Jun 19, 2022
		8252813	Oct 02, 2026	DP U-2028	GAIN	Jun 19, 2027
		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>DESLORATADINE - CLARINEX</u>						
N 021312	001	7618649	Dec 19, 2020	DP U-1017		

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<u>DES Loratadine - Clarinex</u>						
N 021312	002 7618649	Dec 19, 2020	DP U-1017			
<u>DES Loratadine; Pseudoephedrine Sulfate - Clarinex-D 12 Hour</u>						
N 021313	001 6709676	Feb 18, 2021	DP U-707			
	7618649	Dec 19, 2020	DP U-1017			
	8187630	Dec 19, 2020	DP U-1017			
<u>DES Loratadine; Pseudoephedrine Sulfate - Clarinex D 24 Hour</u>						
N 021605	001 6979463	Mar 28, 2022	DP			
	7618649	Dec 19, 2020	DP U-1017			
	7820199	Mar 28, 2022	DP			
<u>DES Mopressin Acetate - NoCDURNA</u>						
N 022517	001 10307459	May 07, 2023	DP		NP	Jun 21, 2021
	7560429	Feb 02, 2024	DP U-2326			
	7947654	Dec 29, 2023	DP			
	8802624	Dec 29, 2023	U-2326			
	9220747	May 07, 2023	U-2326			
	9504647	May 07, 2023	DP U-2326			
	9919025	May 07, 2023	U-2326			
	9974826	Apr 13, 2030	U-2326			
<u>DES Mopressin Acetate - NoCDURNA</u>						
N 022517	002 10137167	May 21, 2029	U-2327		NP	Jun 21, 2021
	10307459	May 07, 2023	DP			
	7560429	Feb 02, 2024	DP U-2326			
	7947654	Dec 29, 2023	DP			
	8802624	Dec 29, 2023	U-2326			
	9220747	May 07, 2023	U-2326			
	9504647	May 07, 2023	DP U-2326			
	9919025	May 07, 2023	U-2326			
	9974826	Apr 13, 2030	U-2327			
<u>DES Mopressin Acetate - NoCTIVA</u>						
N 201656	001 7405203	May 06, 2023	U-1980		NP	Mar 03, 2020
	7579321	May 06, 2023	U-1980			
	7799761	Sep 26, 2024	DP			
	9539302	Jun 15, 2030	DP			
<u>DES Mopressin Acetate - NoCTIVA</u>						
N 201656	002 7405203	May 06, 2023	U-1980		NP	Mar 03, 2020
	7579321	May 06, 2023	U-1980			
	9539302	Jun 15, 2030	DP			
<u>DES Onide - Desonate</u>						
N 021844	001 6387383	Aug 03, 2020	DS DP U-783			
<u>DES Onide - 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>DES Oximetasonone - Topicort</u>						
N 204141	001 8277780	Sep 01, 2028	DP U-1408			
	8715624	May 26, 2026	DP U-1408			
<u>DES Venlafaxine Succinate - Pristiq</u>						
N 021992	001 6673838	Mar 01, 2022	DS U-1364		M-222	Feb 06, 2021
	6673838	Mar 01, 2022	DS U-860			
	8269040	Jul 05, 2027	DS			
<u>DES Venlafaxine Succinate - Pristiq</u>						
N 021992	002 6673838	Mar 01, 2022	DS U-1364		M-222	Feb 06, 2021
	6673838	Mar 01, 2022	DS U-860			
	8269040	Jul 05, 2027	DS			
<u>DES Venlafaxine Succinate - Pristiq</u>						
N 021992	003 6673838	Mar 01, 2022	DS U-1364		M-222	Feb 06, 2021
	6673838	Mar 01, 2022	DS U-860			
	8269040	Jul 05, 2027	DS			

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<u>DEUTETRABENAZINE - AUSTEDO</u>						
N 208082 001	8524733	Mar 27, 2031	DS DP		I-751	Aug 30, 2020
	9233959	Sep 18, 2033	DP		NCE	Apr 03, 2022
	9296739	Sep 18, 2033	DP		ODE-134	Apr 03, 2024
	9550780	Sep 18, 2033	DS DP U-1846			
	9550780	Sep 18, 2033	DS DP U-1995			
	9814708	Sep 18, 2033	DP			
<u>DEUTETRABENAZINE - AUSTEDO</u>						
N 208082 002	8524733	Mar 27, 2031	DS DP		I-751	Aug 30, 2020
	9233959	Sep 18, 2033	DP		NCE	Apr 03, 2022
	9296739	Sep 18, 2033	DP		ODE-134	Apr 03, 2024
	9550780	Sep 18, 2033	DS DP U-1846			
	9550780	Sep 18, 2033	DS DP U-1995			
	9814708	Sep 18, 2033	DP			
<u>DEUTETRABENAZINE - AUSTEDO</u>						
N 208082 003	8524733	Mar 27, 2031	DS DP		I-751	Aug 30, 2020
	9233959	Sep 18, 2033	DP		NCE	Apr 03, 2022
	9296739	Sep 18, 2033	DP		ODE-134	Apr 03, 2024
	9550780	Sep 18, 2033	DS DP U-1846			
	9550780	Sep 18, 2033	DS DP U-1995			
	9814708	Sep 18, 2033	DP			
<u>DEXAMETHASONE - OZURDEX</u>						
N 022315 001	10076526	Jan 09, 2023	DP			
	6726918	Oct 20, 2020	DP U-1204			
	6726918	Oct 20, 2020	DP U-1205			
	6899717	Nov 01, 2023	U-1206			
	7033605	Oct 20, 2020	DP			
	7767223	Nov 28, 2021	DP			
	8034366	Jan 09, 2023	DP U-1204			
	8034366	Jan 09, 2023	DP U-1205			
	8034370	Jan 09, 2023	DP			
	8043628	Oct 20, 2020	U-1205			
	8063031	Oct 20, 2020	DP			
	8088407	Oct 20, 2020	U-1205			
	8506987	Jan 09, 2023	U-1204			
	8506987	Jan 09, 2023	U-1205			
	9012437	Oct 20, 2020	U-1205			
	9192511	Jan 09, 2023	DP			
	9283178	Oct 20, 2020	U-1205			
	9592242	Oct 20, 2020	U-1989			
	9592242	Oct 20, 2020	U-1990			
	9775849	Oct 20, 2020	U-1989			
	9775849	Oct 20, 2020	U-1990			
<u>DEXAMETHASONE - DEXTENZA</u>						
N 208742 001	8409606	May 14, 2030	DP		I-800	Jun 20, 2022
	8563027	Feb 12, 2030	U-2487		NP	Nov 30, 2021
	9254267	Sep 11, 2024	DP			
<u>DEXAMETHASONE - DEXYCU KIT</u>						
N 208912 001	10022502	Sep 12, 2020	U-2340		NP	Feb 09, 2021
	10028965	May 23, 2034	U-2340			
	10159683	May 23, 2034	DP			
	6960346	Jul 03, 2023	DP			
	7560120	Sep 05, 2022	DP			
<u>DEXAMETHASONE - HEMADY</u>						
N 211379 001	10537585	Dec 18, 2037	DP		ODE-271	Oct 03, 2026
<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	6462058	Jun 15, 2020	DS DP U-949			
	6462058	Jun 15, 2020	DS DP U-950			
	6462058	Jun 15, 2020	DS DP U-951			

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<u>DEXLANSOPRAZOLE - DEXILANT</u>						
N 022287 001	6664276	Jan 30, 2023	DS DP U-1507			
	6664276	Jan 30, 2023	DS DP U-949			
	6664276	Jan 30, 2023	DS DP U-950			
	6664276	Jan 30, 2023	DS DP U-951			
	6664276*PED	Jul 30, 2023				
	6939971	Jun 15, 2020		U-949		
	6939971	Jun 15, 2020		U-950		
	6939971	Jun 15, 2020		U-951		
	7285668	Jun 15, 2020	DS			
	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				
	8722084	Oct 15, 2023		DP		
	8722084*PED	Apr 15, 2024				
	8784885	Oct 15, 2023		DP	U-1552	
	8784885	Oct 15, 2023		DP	U-1553	
	8784885	Oct 15, 2023		DP	U-1554	
	8784885*PED	Apr 15, 2024				
	8871273	Jan 11, 2028		DP		
	9011926	Feb 24, 2026		DP		
	9145389	Jun 15, 2020	DS DP			
	9233103	Mar 05, 2032			U-1805	
	9238029	Jan 17, 2026		DP		
<u>DEXLANSOPRAZOLE - DEXILANT</u>						
N 022287 002	6462058	Jun 15, 2020	DS DP U-949			
	6462058	Jun 15, 2020	DS DP U-950			
	6462058	Jun 15, 2020	DS DP U-951			
	6664276	Jan 30, 2023	DS DP U-1507			
	6664276	Jan 30, 2023	DS DP U-949			
	6664276	Jan 30, 2023	DS DP U-950			
	6664276	Jan 30, 2023	DS DP U-951			
	6664276*PED	Jul 30, 2023				
	6939971	Jun 15, 2020		U-949		
	6939971	Jun 15, 2020		U-950		
	6939971	Jun 15, 2020		U-951		
	7285668	Jun 15, 2020	DS			
	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				
	8722084	Oct 15, 2023		DP		
	8722084*PED	Apr 15, 2024				
	8784885	Oct 15, 2023		DP	U-1552	
	8784885	Oct 15, 2023		DP	U-1553	
	8784885	Oct 15, 2023		DP	U-1554	
	8784885*PED	Apr 15, 2024				
	8871273	Jan 11, 2028		DP		
	9011926	Feb 24, 2026		DP		
	9145389	Jun 15, 2020	DS DP			
	9233103	Mar 05, 2032			U-1805	
	9238029	Jan 17, 2026		DP		
<u>DEXLANSOPRAZOLE - DEXILANT SOLUTAB</u>						
N 208056 001	6462058	Jun 15, 2020	DS DP U-950			
	6462058	Jun 15, 2020	DS DP U-951			
	6462058*PED	Dec 15, 2020				
	6664276	Jan 30, 2023	DS DP U-950			
	6664276	Jan 30, 2023	DS DP U-951			
	6664276*PED	Jul 30, 2023				
	6939971	Jun 15, 2020		U-950		
	6939971	Jun 15, 2020		U-951		

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<u>DEXLANSOPRAZOLE - DEXILANT SOLUTAB</u>						
N 208056 001	6939971*PED	Dec 15, 2020				
	7285668	Jun 15, 2020	DS			
	7285668*PED	Dec 15, 2020				
	8461187	Jan 17, 2026	DP			
	8461187*PED	Jul 17, 2026				
	8784885	Oct 15, 2023	DP			
	8784885*PED	Apr 15, 2024				
	8871273	Jan 11, 2028	DP			
	8871273*PED	Jul 11, 2028				
	9011926	Feb 24, 2026	DP			
	9145389	Jun 15, 2020	DS DP			
	9238029	Jan 17, 2026	DP			
	9241910	Mar 10, 2029	DP			
<u>DEXMEDETOMIDINE HYDROCHLORIDE - PRECEDEX</u>						
N 021038 002	10016396	Jan 04, 2032	DP			
	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			
	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			
	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 - DEXMEDETOMIDINE HYDROCHLORIDE</u>						
N 206628 003	9649296	Apr 20, 2036	DP			
	9717796	Apr 20, 2036	DP			
<u>DEXMEDETOMIDINE HYDROCHLORIDE - DEXMEDETOMIDINE HYDROCHLORIDE</u>						
N 206628 004	9649296	Apr 20, 2036	DP			
	9717796	Apr 20, 2036	DP			
<u>DEXRAZOXANE HYDROCHLORIDE - TOTECT</u>						
N 022025 001	6727253	Mar 13, 2020		U-829		

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<u>DEXTROMETHORPHAN HYDROBROMIDE; GUAIFENESIN - MUCINEX DM</u>						
N 021620 001	6372252	Apr 28, 2020	DP			
	6955821	Apr 28, 2020	DP U-685			
	7838032	Apr 28, 2020	DP			
<u>DEXTROMETHORPHAN HYDROBROMIDE; GUAIFENESIN - MUCINEX DM</u>						
N 021620 002	6372252	Apr 28, 2020	DP			
	6955821	Apr 28, 2020	DP U-685			
	7838032	Apr 28, 2020	DP			
<u>DEXTROMETHORPHAN HYDROBROMIDE; QUINIDINE SULFATE - NUEDEXTA</u>						
N 021879 001	7659282	Aug 13, 2026	U-1093			
	8227484	Jul 17, 2023	U-1093			
<u>DIAZEPAM - VALTOCO</u>						
N 211635 001	10265402	May 11, 2025	DP		NP	Jan 10, 2023
	8895546	Mar 27, 2029	DP		ODE-279	Jan 10, 2027
	8927497	Jul 21, 2025	DP U-2727			
	9642913	May 11, 2025	DP			
	9763876	Mar 27, 2029	DP U-2727			
<u>DIAZEPAM - VALTOCO</u>						
N 211635 002	10265402	May 11, 2025	DP		NP	Jan 10, 2023
	8895546	Mar 27, 2029	DP		ODE-279	Jan 10, 2027
	8927497	Jul 21, 2025	DP U-2727			
	9642913	May 11, 2025	DP			
	9763876	Mar 27, 2029	DP U-2727			
<u>DIAZEPAM - VALTOCO</u>						
N 211635 003	10265402	May 11, 2025	DP		NP	Jan 10, 2023
	8895546	Mar 27, 2029	DP		ODE-279	Jan 10, 2027
	8927497	Jul 21, 2025	DP U-2727			
	9642913	May 11, 2025	DP			
	9763876	Mar 27, 2029	DP U-2727			
<u>DIAZOXIDE - DIAZOXIDE</u>						
A 211050 001					CGT	Jun 17, 2020
<u>DICHLORPHENAMIDE - KEVEYIS</u>						
N 011366 002					ODE-96	Aug 07, 2022
<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					NP	Dec 19, 2021
<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			

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<u>DICLOFENAC POTASSIUM - ZIPSOR</u>						
N 022202	001 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>DIGOXIN - DIGOXIN</u>						
A 213000	001				CGT	Apr 01, 2020
<u>DILTIAZEM HYDROCHLORIDE - CARDIZEM LA</u>						
N 021392	001 6923984	Feb 25, 2021	DP			
<u>DILTIAZEM HYDROCHLORIDE - CARDIZEM LA</u>						
N 021392	002 6923984	Feb 25, 2021	DP			
<u>DILTIAZEM HYDROCHLORIDE - CARDIZEM LA</u>						
N 021392	003 6923984	Feb 25, 2021	DP			
<u>DILTIAZEM HYDROCHLORIDE - CARDIZEM LA</u>						
N 021392	004 6923984	Feb 25, 2021	DP			
<u>DILTIAZEM HYDROCHLORIDE - CARDIZEM LA</u>						
N 021392	005 6923984	Feb 25, 2021	DP			
<u>DILTIAZEM HYDROCHLORIDE - CARDIZEM LA</u>						
N 021392	006 6923984	Feb 25, 2021	DP			
<u>DIMETHYL FUMARATE - TECFIDERA</u>						
N 204063	001 7619001	Jun 20, 2020	U-1384			
	8399514	Feb 07, 2028	U-1384			
<u>DIMETHYL FUMARATE - TECFIDERA</u>						
N 204063	002 7619001	Jun 20, 2020	U-1384			
	8399514	Feb 07, 2028	U-1384			

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<u>DIPHENHYDRAMINE CITRATE; IBUPROFEN - ADVIL PM</u>						
N 021394 001	8263647	May 30, 2022	DP			
<u>DIPHENHYDRAMINE HYDROCHLORIDE; IBUPROFEN - ADVIL PM</u>						
N 021393 001	8883849	Jan 17, 2022		U-1618		
	9155718	Jan 17, 2022	DP			
<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	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 002	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	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>DOLUTEGRAVIR SODIUM - TIVICAY</u>						
N 204790 001	8129385	Oct 05, 2027	DS DP		I-758	Nov 21, 2020
	9242986	Dec 08, 2029	DS DP			
<u>DOLUTEGRAVIR SODIUM - TIVICAY</u>						
N 204790 002	8129385	Oct 05, 2027	DS DP		I-758	Nov 21, 2020
	9242986	Dec 08, 2029	DS DP			
<u>DOLUTEGRAVIR SODIUM - TIVICAY</u>						
N 204790 003	8129385	Oct 05, 2027	DS DP		I-758	Nov 21, 2020
	9242986	Dec 08, 2029	DS DP			
<u>DOLUTEGRAVIR SODIUM; LAMIVUDINE - DOVATO</u>						
N 211994 001	8129385	Oct 05, 2027	DS DP		NC	Apr 08, 2022
	9242986	Dec 08, 2029	DS DP			
<u>DOLUTEGRAVIR SODIUM; RILPIVIRINE HYDROCHLORIDE - JULUCA</u>						
N 210192 001	10426780	Jan 24, 2031	DS DP	U-257	NC	Nov 21, 2020
	6838464	Feb 26, 2021	DS DP			
	7125879	Apr 21, 2025	DS DP	U-257		
	8080551	Apr 11, 2023	DS DP			
	8101629	Aug 09, 2022	DP			
	8129385	Oct 05, 2027	DS DP			
	9242986	Dec 08, 2029	DS DP			

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DONEPEZIL HYDROCHLORIDE - ARICEPT

N 022568	001	8481565	Oct 04, 2026	DP		
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DONEPEZIL HYDROCHLORIDE; MEMANTINE HYDROCHLORIDE - NAMZARIC

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		
		8168209*PED	May 22, 2026			
		8173708	Nov 22, 2025		U-1641	
		8173708*PED	May 22, 2026			
		8283379	Nov 22, 2025		U-1641	
		8283379*PED	May 22, 2026			
		8293794	Nov 22, 2025	DP		
		8329752	Nov 22, 2025	DP		
		8329752*PED	May 22, 2026			
		8338485	Nov 22, 2025	DP		
		8338486	Nov 22, 2025		U-1641	
		8362085	Nov 22, 2025		U-1641	
		8362085*PED	May 22, 2026			
		8580858	Nov 22, 2025		U-1641	
		8598233	Nov 22, 2025	DP		
		8598233*PED	May 22, 2026			

DONEPEZIL HYDROCHLORIDE; MEMANTINE HYDROCHLORIDE - NAMZARIC

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		
		8168209*PED	May 22, 2026			
		8173708	Nov 22, 2025		U-1641	
		8173708*PED	May 22, 2026			
		8283379	Nov 22, 2025		U-1641	
		8283379*PED	May 22, 2026			
		8293794	Nov 22, 2025	DP		
		8329752	Nov 22, 2025	DP		
		8329752*PED	May 22, 2026			
		8338485	Nov 22, 2025	DP		
		8338486	Nov 22, 2025		U-1641	
		8362085	Nov 22, 2025		U-1641	
		8362085*PED	May 22, 2026			
		8580858	Nov 22, 2025		U-1641	
		8598233	Nov 22, 2025	DP		
		8598233*PED	May 22, 2026			

DONEPEZIL HYDROCHLORIDE; MEMANTINE HYDROCHLORIDE - NAMZARIC

N 206439	003	8039009	Mar 24, 2029		U-1641	
		8039009*PED	Sep 24, 2029			
		8058291	Dec 05, 2029		U-1641	
		8168209	Nov 22, 2025	DP		
		8168209*PED	May 22, 2026			
		8173708	Nov 22, 2025		U-1641	
		8173708*PED	May 22, 2026			
		8283379	Nov 22, 2025		U-1641	
		8283379*PED	May 22, 2026			
		8293794	Nov 22, 2025	DP		
		8329752	Nov 22, 2025	DP		
		8329752*PED	May 22, 2026			
		8338485	Nov 22, 2025	DP		
		8338486	Nov 22, 2025		U-1641	
		8362085	Nov 22, 2025		U-1641	
		8362085*PED	May 22, 2026			
		8580858	Nov 22, 2025		U-1641	
		8598233	Nov 22, 2025	DP		
		8598233*PED	May 22, 2026			

DONEPEZIL HYDROCHLORIDE; MEMANTINE HYDROCHLORIDE - NAMZARIC

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		

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<u>DONEPEZIL HYDROCHLORIDE; MEMANTINE HYDROCHLORIDE - NAMZARIC</u>						
N 206439 004	8168209*PED	May 22, 2026				
	8173708	Nov 22, 2025	U-1641			
	8173708*PED	May 22, 2026				
	8283379	Nov 22, 2025	U-1641			
	8283379*PED	May 22, 2026				
	8293794	Nov 22, 2025	DP			
	8329752	Nov 22, 2025	DP			
	8329752*PED	May 22, 2026				
	8338485	Nov 22, 2025	DP			
	8338486	Nov 22, 2025	U-1641			
	8362085	Nov 22, 2025	U-1641			
	8362085*PED	May 22, 2026				
	8580858	Nov 22, 2025	U-1641			
	8598233	Nov 22, 2025	DP			
	8598233*PED	May 22, 2026				
<u>DORAVIRINE - PIFELTRO</u>						
N 210806 001	8486975	Oct 07, 2031	DS DP U-2394		NCE	Aug 30, 2023
	8486975	Oct 07, 2031	DS DP U-2630			
<u>DORAVIRINE; LAMIVUDINE; TENOFOVIR DISOPROXIL FUMARATE - DELSTRIGO</u>						
N 210807 001	8486975	Oct 07, 2031	DS DP U-2395		I-806	Sep 19, 2022
	8486975	Oct 07, 2031	DS DP U-2629		NCE	Aug 30, 2023
<u>DORIPENEM - DORIBAX</u>						
N 022106 001	8247402	Mar 30, 2021	DS DP			
<u>DORIPENEM - DORIBAX</u>						
N 022106 002	8247402	Mar 30, 2021	DS DP			
<u>DOXEPIH HYDROCHLORIDE - SILENOR</u>						
N 022036 001	10238620	May 18, 2027	U-620			
	10548871	Apr 11, 2028	U-620			
	6211229	Feb 17, 2020	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>DOXEPIH HYDROCHLORIDE - SILENOR</u>						
N 022036 002	10238620	May 18, 2027	U-620			
	10548871	Apr 11, 2028	U-620			
	6211229	Feb 17, 2020	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	10058564	Apr 05, 2022	U-1063			
	7211267	Apr 05, 2022	U-925			
	7232572	Apr 05, 2022	U-925			
	7749532	Dec 19, 2027	DP U-1063			
	8206740	Dec 24, 2025	DP U-925			
	8394405	Apr 07, 2024	DP U-925			
	8394406	Apr 07, 2024	DP U-925			
	8470364	Apr 07, 2024	DP U-925			
	8603506	Apr 05, 2022	U-1063			
	8709478	Apr 07, 2024	U-1063			
	9241946	Apr 05, 2022	U-1063			

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<u>DOXYCYCLINE HYCLATE - DORYX</u>						
N 050795 001	6958161	Dec 15, 2022	DP U-918			
	8715724	Feb 03, 2028	DP			
<u>DOXYCYCLINE HYCLATE - DORYX</u>						
N 050795 002	6958161	Dec 15, 2022	DP U-918			
	8715724	Feb 03, 2028	DP			
<u>DOXYCYCLINE HYCLATE - DORYX</u>						
N 050795 003	6958161	Dec 15, 2022	DP U-918			
	8715724	Feb 03, 2028	DP			
<u>DOXYCYCLINE HYCLATE - DORYX</u>						
N 050795 004	6958161	Dec 15, 2022	DP U-918			
	8715724	Feb 03, 2028	DP			
<u>DOXYCYCLINE HYCLATE - DORYX</u>						
N 050795 005	6958161	Dec 15, 2022	DP U-918			
	8715724	Feb 03, 2028	DP			
<u>DOXYCYCLINE HYCLATE - DORYX</u>						
N 050795 006	6958161	Dec 15, 2022	DP U-918			
	8715724	Feb 03, 2028	DP			
<u>DOXYCYCLINE HYCLATE - DORYX MPC</u>						
N 050795 007	6958161	Dec 15, 2022	DP U-918			
	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	6958161	Dec 15, 2022	DP U-918			
	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 - DICLEGIS</u>						
N 021876 001	6340695	Jun 21, 2021	DP U-1382			
<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			
	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		NP	May 23, 2022
	9603860	Jun 28, 2031	U-2553			
<u>DROSPIRENONE; ESTRADIOL - ANGELIO</u>						
N 021355 001	8906890	Oct 22, 2031	DP			
<u>DROSPIRENONE; ETHINYL ESTRADIOL - YASMIN</u>						
N 021098 001	6787531	Aug 31, 2020	DP			

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<u>DROSPIRENONE; ETHINYL ESTRADIOL - YAZ</u>						
N 021676	001	6787531	Aug 31, 2020	DP		
		6958326	Dec 20, 2021	DP		
		7163931	Dec 20, 2021	U-1		
<u>DROSPIRENONE; ETHINYL ESTRADIOL; LEVOMEFOLATE CALCIUM - BEYAZ</u>						
N 022532	001	6441168	Jul 30, 2022	DS		
		6958326	Dec 20, 2021	DP		
		7163931	Mar 03, 2022	U-1		
		8617597	Feb 08, 2030	DP		
<u>DROSPIRENONE; ETHINYL ESTRADIOL; LEVOMEFOLATE CALCIUM - SAFYRAL</u>						
N 022574	001	6441168	Apr 17, 2020	DS		
		6958326	Dec 20, 2021	DP		
		7163931	Mar 03, 2022	U-1		
		8617597	Feb 08, 2030	DP		
<u>DROXIDOPA - NORTHERA</u>						
N 203202	001				ODE-61	Feb 18, 2021
<u>DROXIDOPA - NORTHERA</u>						
N 203202	002				ODE-61	Feb 18, 2021
<u>DROXIDOPA - NORTHERA</u>						
N 203202	003				ODE-61	Feb 18, 2021
<u>DULOXETINE HYDROCHLORIDE - DRIZALMA SPRINKLE</u>						
N 212516	001	10413525	Apr 13, 2037	DP		
		9839626	Apr 13, 2037	DP		
<u>DULOXETINE HYDROCHLORIDE - DRIZALMA SPRINKLE</u>						
N 212516	002	10413525	Apr 13, 2037	DP		
		9839626	Apr 13, 2037	DP		
<u>DULOXETINE HYDROCHLORIDE - DRIZALMA SPRINKLE</u>						
N 212516	003	10413525	Apr 13, 2037	DP		
		9839626	Apr 13, 2037	DP		
<u>DULOXETINE HYDROCHLORIDE - DRIZALMA SPRINKLE</u>						
N 212516	004	10413525	Apr 13, 2037	DP		
		9839626	Apr 13, 2037	DP		
<u>DUVELISIB - COPIKTRA</u>						
N 211155	001	8193182	Feb 13, 2030	DS	NCE	Sep 24, 2023
		9216982	Jan 05, 2029	U-2412	ODE-208	Sep 24, 2025
		9216982	Jan 05, 2029	U-2413	ODE-209	Sep 24, 2025
		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	8193182	Feb 13, 2030	DS	NCE	Sep 24, 2023
		9216982	Jan 05, 2029	U-2412	ODE-208	Sep 24, 2025
		9216982	Jan 05, 2029	U-2413	ODE-209	Sep 24, 2025
		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</u>						
N 209176	001	6933310	Nov 13, 2020	U-2013	NCE	May 05, 2022
					ODE-144	May 05, 2024
<u>EDARAVONE - RADICAVA</u>						
N 209176	002	6933310	Nov 13, 2020	U-2013	NCE	May 05, 2022
<u>EDOXYBAN TOSYLATE - SAVAYSA</u>						
N 206316	001	7365205	Jun 12, 2023	DS	M-243	Aug 09, 2022
		9149532	Mar 28, 2028	DP	NCE	Jan 08, 2020

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<u>EDOXABAN TOSYLATE - SAVAYSA</u>						
N 206316 002	7365205	Jun 12, 2023	DS		M-243	Aug 09, 2022
	9149532	Mar 28, 2028	DP		NCE	Jan 08, 2020
<u>EDOXABAN TOSYLATE - SAVAYSA</u>						
N 206316 003	7365205	Jun 12, 2023	DS		M-243	Aug 09, 2022
	9149532	Mar 28, 2028	DP		NCE	Jan 08, 2020
<u>EMTRICITABINE; EMTRICITABINE; TENOFOVIR DISOPROXIL FUMARATE - ATRIPLA</u>						
N 021937 001	6642245	Nov 04, 2020		U-1170		
	6642245	Nov 04, 2020		U-750		
	6703396	Mar 09, 2021	DS DP			
	8592397	Jan 13, 2024		DP U-1170		
	8592397	Jan 13, 2024		DP U-750		
	8598185	Apr 28, 2029		DP		
	8716264	Jan 13, 2024		DP U-257		
	9018192	Jun 13, 2026		U-1170		
	9018192	Jun 13, 2026		U-750		
	9457036	Jan 13, 2024		DP U-257		
	9545414	Jun 13, 2026		DP U-1170		
	9545414	Jun 13, 2026		DP U-750		
	9744181	Jan 13, 2024		DP U-257		
<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		
	7214506	Oct 05, 2021		U-281		
	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>ELAGOLIX SODIUM - ORILISSA</u>						
N 210450 001	10537572	Sep 01, 2036		U-2735	NCE	Jul 23, 2023
	6872728	Jan 25, 2021	DS DP			
	7056927	Sep 10, 2024	DS DP			
	7176211	Jul 06, 2024		U-2360		
	7179815	Mar 07, 2021		U-2360		
	7419983	Jul 06, 2024	DS DP	U-2360		
	7462625	Jan 25, 2021	DS DP	U-2360		
<u>ELAGOLIX SODIUM - ORILISSA</u>						
N 210450 002	6872728	Jan 25, 2021	DS DP		NCE	Jul 23, 2023
	7056927	Sep 10, 2024	DS DP			
	7176211	Jul 06, 2024		U-2360		
	7179815	Mar 07, 2021		U-2360		
	7419983	Jul 06, 2024	DS DP	U-2360		
	7462625	Jan 25, 2021	DS DP	U-2360		
<u>ELBASVIR; GRAZOPREXIVIR - ZEPATIER</u>						
N 208261 001	7973040	Jul 24, 2029	DS DP	U-1813	NCE	Jan 28, 2021
	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	NCE	Feb 12, 2023
	10081621	Mar 25, 2031		DP U-2652	NCE	Oct 21, 2024
	10239867	Apr 09, 2027	DS DP	U-2653	ODE-275	Oct 21, 2026
	7495103	May 20, 2027	DS DP			
	7645789	May 01, 2027	DS DP			
	7776905	Jun 03, 2027	DS DP			
	8324242	Aug 05, 2027		U-2645		
	8354427	Jul 06, 2026		U-2646		
	8410274	Dec 28, 2026		DP		
	8415387	Nov 12, 2027		U-2645		
	8598181	May 01, 2027		U-2645		
	8623905	May 01, 2027	DS DP			

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<u>ELEXACAFTOR, IVACAFTOR, TEZACAFTOR; IVACAFTOR - TRIKAFTA (COPACKAGED)</u>						
N 212273 001	8629162	Jun 24, 2025			U-2648	
	8754224	Dec 28, 2026	DS DP			
	9012496	Jul 15, 2033			U-2649	
	9670163	Dec 28, 2026		DP	U-2650	
	9931334	Dec 28, 2026		DP	U-2650	
	9974781	Apr 09, 2027		DP	U-2645	
<u>ELIGLUSTAT TARTRATE - CERDELGA</u>						
N 205494 001	6916802	Apr 29, 2022	DS	U-1571		Aug 19, 2021
	7196205	Jun 26, 2026	DS		ODE-73	
	7253185	Apr 29, 2022		DP		
	7615573	Apr 29, 2022			U-1571	
<u>ELTROMBOPAG OLAMINE - PROMACTA</u>						
N 022291 001	7160870	Nov 20, 2022	DS DP	U-1306	ODE-210	Nov 16, 2025
	7160870	Nov 20, 2022	DS DP	U-2451	ODE-75	Aug 26, 2021
	7160870*PED	May 20, 2023			PED	Feb 26, 2022
	7332481	May 24, 2021			U-1306	
	7332481	May 24, 2021			U-2451	
	7332481*PED	Nov 24, 2021				
	7452874	May 24, 2021	DS DP			
	7452874*PED	Nov 24, 2021				
	7473686	May 24, 2021	DS DP	U-1306		
	7473686	May 24, 2021	DS DP	U-2451		
	7473686*PED	Nov 24, 2021				
	7547719	Jul 13, 2025	DS DP	U-1306		
	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				
	7790704	May 24, 2021			U-1306	
	7790704	May 24, 2021			U-2451	
	7790704*PED	Nov 24, 2021				
	7795293	May 21, 2023			U-1306	
	7795293	May 21, 2023			U-2451	
	7795293*PED	Nov 21, 2023				
	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		
	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	7160870	Nov 20, 2022	DS DP	U-1306	ODE-210	Nov 16, 2025
	7160870	Nov 20, 2022	DS DP	U-2451	ODE-75	Aug 26, 2021
	7160870*PED	May 20, 2023			PED	Feb 26, 2022
	7332481	May 24, 2021			U-1306	
	7332481	May 24, 2021			U-2451	
	7332481*PED	Nov 24, 2021				
	7452874	May 24, 2021	DS DP			
	7452874*PED	Nov 24, 2021				
	7473686	May 24, 2021	DS DP	U-1306		
	7473686	May 24, 2021	DS DP	U-2451		
	7473686*PED	Nov 24, 2021				
	7547719	Jul 13, 2025	DS DP	U-1306		
	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				
	7790704	May 24, 2021			U-1306	

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<u>ELTROMBOPAG OLAMINE - PROMACTA</u>						
N 022291 002	7790704	May 24, 2021	U-2451			
	7790704*PED	Nov 24, 2021				
	7795293	May 21, 2023	U-1306			
	7795293	May 21, 2023	U-2451			
	7795293	May 21, 2023	U-930			
	7795293*PED	Nov 21, 2023				
	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	7160870	Nov 20, 2022	DS DP U-1306		ODE-210	Nov 16, 2025
	7160870	Nov 20, 2022	DS DP U-2451		ODE-75	Aug 26, 2021
	7160870*PED	May 20, 2023			PED	Feb 26, 2022
	7332481	May 24, 2021	U-1306			
	7332481	May 24, 2021	U-2451			
	7332481*PED	Nov 24, 2021				
	7452874	May 24, 2021	DS DP			
	7452874*PED	Nov 24, 2021				
	7473686	May 24, 2021	DS DP U-1306			
	7473686	May 24, 2021	DS DP U-2451			
	7473686*PED	Nov 24, 2021				
	7547719	Jul 13, 2025	DS DP U-1306			
	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				
	7790704	May 24, 2021	U-1306			
	7790704	May 24, 2021	U-2451			
	7790704*PED	Nov 24, 2021				
	7795293	May 21, 2023	U-1306			
	7795293	May 21, 2023	U-2451			
	7795293	May 21, 2023	U-930			
	7795293*PED	Nov 21, 2023				
	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			
	8828430*PED	Feb 01, 2028				
<u>ELTROMBOPAG OLAMINE - PROMACTA</u>						
N 022291 004	7160870	Nov 20, 2022	DS DP U-1306		ODE-210	Nov 16, 2025
	7160870	Nov 20, 2022	DS DP U-2451		ODE-75	Aug 26, 2021
	7160870*PED	May 20, 2023			PED	Feb 26, 2022
	7332481	May 24, 2021	U-1306			
	7332481	May 24, 2021	U-2451			
	7332481*PED	Nov 24, 2021				
	7452874	May 24, 2021	DS DP			
	7452874*PED	Nov 24, 2021				
	7473686	May 24, 2021	DS DP U-1306			

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<u>ELTROMBOPAG OLAMINE - PROMACTA</u>						
N 022291 004	7473686	May 24, 2021	DS DP U-2451			
	7473686*PED	Nov 24, 2021				
	7547719	Jul 13, 2025	DS DP U-1306			
	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				
	7790704	May 24, 2021		U-1306		
	7790704	May 24, 2021		U-2451		
	7790704*PED	Nov 24, 2021				
	7795293	May 21, 2023		U-1306		
	7795293	May 21, 2023		U-2451		
	7795293*PED	Nov 21, 2023				
	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	7160870	Nov 20, 2022	DS DP U-1306		ODE-210	Nov 16, 2025
	7160870	Nov 20, 2022	DS DP U-1575		ODE-75	Aug 26, 2021
	7160870	Nov 20, 2022	DS DP U-1714		PED	Feb 26, 2022
	7160870	Nov 20, 2022	DS DP U-930			
	7160870*PED	May 20, 2023				
	7332481	May 24, 2021		U-1306		
	7332481	May 24, 2021		U-1575		
	7332481	May 24, 2021		U-1714		
	7332481	May 24, 2021		U-930		
	7332481*PED	Nov 24, 2021				
	7452874	May 24, 2021	DS DP U-1714			
	7452874*PED	Nov 24, 2021				
	7473686	May 24, 2021	DS DP U-1306			
	7473686	May 24, 2021	DS DP U-1575			
	7473686	May 24, 2021	DS DP U-1714			
	7473686	May 24, 2021	DS DP U-930			
	7473686*PED	Nov 24, 2021				
	7547719	Jul 13, 2025	DS DP U-1306			
	7547719	Jul 13, 2025	DS DP U-1575			
	7547719	Jul 13, 2025	DS DP U-930			
	7547719*PED	Jan 13, 2026				
	7790704	May 24, 2021		U-1306		
	7790704	May 24, 2021		U-1575		
	7790704	May 24, 2021		U-930		
	7790704*PED	Nov 24, 2021				
	7795293	May 21, 2023		U-1306		
	7795293	May 21, 2023		U-1575		
	7795293	May 21, 2023		U-930		
	7795293*PED	Nov 21, 2023				
<u>ELTROMBOPAG OLAMINE - PROMACTA KIT</u>						
N 207027 001	7160870	Nov 20, 2022	DS DP U-1306		ODE-74	Aug 26, 2021
	7160870	Nov 20, 2022	DS DP U-1736		PED	Feb 26, 2022
	7160870*PED	May 20, 2023				
	7332481	May 24, 2021		U-1306		
	7332481	May 24, 2021		U-1736		
	7332481*PED	Nov 24, 2021				
	7452874	May 24, 2021	DS DP			
	7452874*PED	Nov 24, 2021				
	7473686	May 24, 2021	DS DP U-1306			
	7473686	May 24, 2021	DS DP U-1736			

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<u>ELTROMBOPAG OLAMINE - PROMACTA KIT</u>						
N 207027 001	7473686*PED	Nov 24, 2021				
	7547719	Jul 13, 2025	DS DP U-1306			
	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				
	7790704	May 24, 2021	U-1306			
	7790704	May 24, 2021	U-1736			
	7790704*PED	Nov 24, 2021				
	7795293	May 21, 2023	U-1306			
	7795293	May 21, 2023	U-1736			
	7795293*PED	Nov 24, 2023				
<u>ELTROMBOPAG OLAMINE - PROMACTA KIT</u>						
N 207027 002	7160870	Nov 20, 2022	DS DP U-1306			
	7160870	Nov 20, 2022	DS DP U-1736			
	7160870*PED	May 20, 2023				
	7332481	May 24, 2021	U-1306			
	7332481	May 24, 2021	U-1736			
	7332481*PED	Nov 24, 2021				
	7452874	May 24, 2021	DS DP			
	7452874*PED	Nov 24, 2021				
	7473686	May 24, 2021	DS DP U-1306			
	7473686	May 24, 2021	DS DP U-1736			
	7473686*PED	Nov 24, 2021				
	7547719	Jul 13, 2025	DS DP U-1306			
	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				
	7790704	May 24, 2021	U-1306			
	7790704	May 24, 2021	U-1736			
	7790704*PED	Nov 24, 2021				
	7795293	May 21, 2023	U-1306			
	7795293	May 21, 2023	U-1736			
	7795293*PED	Nov 21, 2023				
<u>ELUXADOLINE - VIBERZI</u>						
N 206940 001	10188632	Mar 14, 2033	DP		NCE	May 27, 2020
	10213415	Mar 14, 2025	DS U-2152			
	7741356	Mar 25, 2028	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>ELUXADOLINE - VIBERZI</u>						
N 206940 002	10188632	Mar 14, 2033	DP		NCE	May 27, 2020
	10213415	Mar 14, 2025	DS U-2152			
	7741356	Mar 25, 2028	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			

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<u>ELUXADOLINE - VIBERZI</u>						
N 206940 002	10188632	Mar 14, 2033	DP		NCE	May 27, 2020
	10213415	Mar 14, 2025	DS	U-2152		
	7741356	Mar 25, 2028	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	
	7635704	Oct 26, 2026	DS	DP	U-257	
	8981103	Oct 26, 2026	DS	DP		
<u>ELVITEGRAVIR - VITEKTA</u>						
N 203093 002	7176220	Aug 27, 2026	DS	DP	U-257	
	7635704	Oct 26, 2026	DS	DP	U-257	
	8981103	Oct 26, 2026	DS	DP		
<u>EMPAGLIFLOZIN - JARDIANCE</u>						
N 204629 001	10258637	Apr 03, 2034			U-2290	
	7579449	Nov 05, 2025	DS			
	7713938	Apr 15, 2027	DS	DP		
	8551957	Oct 14, 2029			U-1651	
	9949997	May 17, 2034			U-2292	
	9949998	Jun 11, 2034			U-2290	
<u>EMPAGLIFLOZIN - JARDIANCE</u>						
N 204629 002	10258637	Apr 03, 2034			U-2290	
	7579449	Nov 05, 2025	DS			
	7713938	Apr 15, 2027	DS	DP		
	8551957	Oct 14, 2029			U-1651	
	9949997	May 17, 2034			U-2292	
	9949998	Jun 11, 2034			U-2290	
<u>EMPAGLIFLOZIN; LINAGLIPTIN - GLYXAMBI</u>						
N 206073 001	10258637	Apr 03, 2034			U-2290	
	7407955	May 02, 2025	DS	DP		
	7579449	Nov 05, 2025	DS			
	7713938	Apr 15, 2027	DS	DP		
	8119648	Aug 12, 2023			U-1651	
	8178541	Aug 12, 2023	DP		U-1653	
	8178541	Aug 12, 2023	DP		U-1654	
	8551957	Oct 14, 2029	DP		U-1651	
	8673927	May 04, 2027			U-1652	
	8883805	Nov 26, 2025	DP			
	9173859	May 04, 2027	DP		U-1772	
	9949998	Jun 11, 2034			U-2290	
<u>EMPAGLIFLOZIN; LINAGLIPTIN - GLYXAMBI</u>						
N 206073 002	10258637	Apr 03, 2034			U-2290	
	7407955	May 02, 2025	DS	DP		
	7579449	Nov 05, 2025	DS			
	7713938	Apr 15, 2027	DS	DP		
	8119648	Aug 12, 2023			U-1651	
	8178541	Aug 12, 2023	DP		U-1653	
	8178541	Aug 12, 2023	DP		U-1654	
	8551957	Oct 14, 2029	DP		U-1651	
	8673927	May 04, 2027			U-1652	
	8883805	Nov 26, 2025	DP			
	9173859	May 04, 2027	DP		U-1772	
	9949998	Jun 11, 2034			U-2290	

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<u>EMPAGLIFLOZIN; LINAGLIPTIN; METFORMIN HYDROCHLORIDE - TRIJARDY XR</u>						
N 212614 001	10022379	Apr 02, 2029	DP U-2732			
	10258637	Apr 03, 2034	U-2731			
	10406172	Jun 15, 2030	DP U-2733			
	6488962	Jun 20, 2020	DP			
	7407955	May 02, 2025	DS DP			
	7579449	Nov 05, 2025	DS			
	7713938	Apr 15, 2027	DS DP			
	8119648	Aug 12, 2023	U-1652			
	8178541	Aug 12, 2023	DP U-1652			
	8551957	Oct 14, 2029	DP U-2730			
	8673927	May 04, 2027	U-1652			
	8883805	Nov 26, 2025	DP			
	9155705	May 21, 2030	DP			
	9173859	May 04, 2027	DP U-1652			
	9173859	May 04, 2027	DP U-2730			
	9415016	Apr 02, 2029	DP			
	9949998	Jun 11, 2034	U-2731			
<u>EMPAGLIFLOZIN; LINAGLIPTIN; METFORMIN HYDROCHLORIDE - TRIJARDY XR</u>						
N 212614 002	10022379	Apr 02, 2029	DP U-2732			
	10258637	Apr 03, 2034	U-2731			
	10406172	Jun 15, 2030	DP U-2733			
	6488962	Jun 20, 2020	DP			
	7407955	May 02, 2025	DS DP			
	7579449	Nov 05, 2025	DS			
	7713938	Apr 15, 2027	DS DP			
	8119648	Aug 12, 2023	U-1652			
	8178541	Aug 12, 2023	DP U-1652			
	8551957	Oct 14, 2029	DP U-2730			
	8673927	May 04, 2027	U-1652			
	8883805	Nov 26, 2025	DP			
	9155705	May 21, 2030	DP			
	9173859	May 04, 2027	DP U-1652			
	9173859	May 04, 2027	DP U-2730			
	9415016	Apr 02, 2029	DP			
	9949998	Jun 11, 2034	U-2731			
<u>EMPAGLIFLOZIN; LINAGLIPTIN; METFORMIN HYDROCHLORIDE - TRIJARDY XR</u>						
N 212614 003	10022379	Apr 02, 2029	DP U-2732			
	10258637	Apr 03, 2034	U-2731			
	10406172	Jun 15, 2030	DP U-2733			
	6488962	Jun 20, 2020	DP			
	7407955	May 02, 2025	DS DP			
	7579449	Nov 05, 2025	DS			
	7713938	Apr 15, 2027	DS DP			
	8119648	Aug 12, 2023	U-1652			
	8178541	Aug 12, 2023	DP U-1652			
	8551957	Oct 14, 2029	DP U-2730			
	8673927	May 04, 2027	U-1652			
	8883805	Nov 26, 2025	DP			
	9155705	May 21, 2030	DP			
	9173859	May 04, 2027	DP U-1652			
	9173859	May 04, 2027	DP U-2730			
	9415016	Apr 02, 2029	DP			
	9949998	Jun 11, 2034	U-2731			
<u>EMPAGLIFLOZIN; LINAGLIPTIN; METFORMIN HYDROCHLORIDE - TRIJARDY XR</u>						
N 212614 004	10022379	Apr 02, 2029	DP U-2732			
	10258637	Apr 03, 2034	U-2731			
	10406172	Jun 15, 2030	DP U-2733			
	6488962	Jun 20, 2020	DP			
	7407955	May 02, 2025	DS DP			
	7579449	Nov 05, 2025	DS			
	7713938	Apr 15, 2027	DS DP			
	8119648	Aug 12, 2023	U-1652			
	8178541	Aug 12, 2023	DP U-1652			
	8551957	Oct 14, 2029	DP U-2730			
	8673927	May 04, 2027	U-1652			
	9155705	May 21, 2030	DP			

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<u>EMPAGLIFLOZIN; LINAGLIPTIN; METFORMIN HYDROCHLORIDE - TRIJARDY XR</u>						
N 212614	004	9173859	May 04, 2027	DP	U-1652	
		9173859	May 04, 2027	DP	U-2730	
		9415016	Apr 02, 2029	DP		
		9949998	Jun 11, 2034		U-2731	
<u>EMPAGLIFLOZIN; METFORMIN HYDROCHLORIDE - SYNJARDY</u>						
N 206111	001	10258637	Apr 03, 2034		U-2290	
		7579449	Nov 05, 2025	DS		
		7713938	Apr 15, 2027	DS	DP	
<u>EMPAGLIFLOZIN; METFORMIN HYDROCHLORIDE - SYNJARDY</u>						
N 206111	002	10258637	Apr 03, 2034		U-2290	
		7579449	Nov 05, 2025	DS		
		7713938	Apr 15, 2027	DS	DP	
<u>EMPAGLIFLOZIN; METFORMIN HYDROCHLORIDE - SYNJARDY</u>						
N 206111	003	10258637	Apr 03, 2034		U-2290	
		7579449	Nov 05, 2025	DS		
		7713938	Apr 15, 2027	DS	DP	
<u>EMPAGLIFLOZIN; METFORMIN HYDROCHLORIDE - SYNJARDY</u>						
N 206111	004	10258637	Apr 03, 2034		U-2290	
		7579449	Nov 05, 2025	DS		
		7713938	Apr 15, 2027	DS	DP	
<u>EMPAGLIFLOZIN; METFORMIN HYDROCHLORIDE - SYNJARDY XR</u>						
N 208658	001	10258637	Apr 03, 2034		U-2290	
		6488962	Jun 20, 2020	DP		
		7579449	Nov 05, 2025	DS		
		7713938	Apr 15, 2027	DS	DP	
		9949998	Jun 11, 2034		U-2290	
<u>EMPAGLIFLOZIN; METFORMIN HYDROCHLORIDE - SYNJARDY XR</u>						
N 208658	002	10258637	Apr 03, 2034		U-2290	
		6488962	Jun 20, 2020	DP		
		7579449	Nov 05, 2025	DS		
		7713938	Apr 15, 2027	DS	DP	
		9949998	Jun 11, 2034		U-2290	
<u>EMPAGLIFLOZIN; METFORMIN HYDROCHLORIDE - SYNJARDY XR</u>						
N 208658	003	10258637	Apr 03, 2034		U-2290	
		6488962	Jun 20, 2020	DP		
		7579449	Nov 05, 2025	DS		
		7713938	Apr 15, 2027	DS	DP	
		9949998	Jun 11, 2034		U-2290	
<u>EMPAGLIFLOZIN; METFORMIN HYDROCHLORIDE - SYNJARDY XR</u>						
N 208658	004	10258637	Apr 03, 2034		U-2290	
		6488962	Jun 20, 2020	DP		
		7579449	Nov 05, 2025	DS		
		7713938	Apr 15, 2027	DS	DP	
		9949998	Jun 11, 2034		U-2290	
<u>EMTRICITABINE - EMTRIVA</u>						
N 021500	001	6642245	Nov 04, 2020		U-257	
		6642245	Nov 04, 2020		U-541	
		6703396	Mar 09, 2021	DS	DP	
<u>EMTRICITABINE - EMTRIVA</u>						
N 021896	001	6642245	Nov 04, 2020		U-257	
		6703396	Mar 09, 2021	DS	DP	
<u>EMTRICITABINE; RILPIVIRINE HYDROCHLORIDE; TENOFOVIR ALAFENAMIDE FUMARATE - ODEFSEY</u>						
N 208351	001	6642245	Nov 04, 2020		U-257	M-206 Aug 21, 2020
		6642245*PED	May 04, 2021			M-207 Aug 21, 2020
		6703396	Mar 09, 2021	DS	DP	NCE Nov 05, 2020
		6703396*PED	Sep 09, 2021			
		6838464	Feb 26, 2021	DS	DP	
		7125879	Apr 21, 2025	DS	DP	U-257
		7390791	May 07, 2022	DS	DP	

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EMTRICITABINE; RILPIVIRINE HYDROCHLORIDE; TENOFOVIR ALAFENAMIDE FUMARATE - ODEFSEY

N 208351	001	7803788	Feb 02, 2022			U-257
		8080551	Apr 11, 2023	DS DP		
		8101629	Aug 09, 2022	DP		
		8754065	Aug 15, 2032	DS DP		U-257
		9296769	Aug 15, 2032	DS DP		U-257

EMTRICITABINE; RILPIVIRINE HYDROCHLORIDE; TENOFOVIR DISOPROXIL FUMARATE - COMPLERA

N 202123	001	6642245	Nov 04, 2020			U-257
		6703396	Mar 09, 2021	DS DP		
		6838464	Feb 26, 2021	DS DP		
		7125879	Apr 21, 2025	DS DP		U-257
		8080551	Apr 11, 2023	DS DP		
		8101629	Aug 09, 2022	DP		
		8592397	Jan 13, 2024	DP		U-257
		8716264	Jan 13, 2024	DP		U-257
		8841310	Dec 09, 2025	DP		U-257
		9457036	Jan 13, 2024	DP		U-257
		9744181	Jan 13, 2024	DP		U-257

EMTRICITABINE; TENOFOVIR ALAFENAMIDE FUMARATE - DESCOVY

N 208215	001	6642245	Nov 04, 2020			U-1663		
		6642245*PED	May 04, 2021				I-812	Oct 03, 2022
		6703396	Mar 09, 2021	DS DP			NCE	Nov 05, 2020
		6703396*PED	Sep 09, 2021				NPP	Sep 25, 2020
		7390791	May 07, 2022	DS DP			ODE-284	Sep 28, 2024
		7803788	Feb 02, 2022			U-1663	ODE-285	Sep 28, 2024
		8754065	Aug 15, 2032	DS DP		U-1259		
		8754065	Aug 15, 2032	DS DP		U-1663		
		8754065	Aug 15, 2032	DS DP		U-257		
		9296769	Aug 15, 2032	DS DP		U-1259		
		9296769	Aug 15, 2032	DS DP		U-1663		
		9296769	Aug 15, 2032	DS DP		U-257		

EMTRICITABINE; TENOFOVIR DISOPROXIL FUMARATE - TRUVADA

N 021752	001	6642245	Nov 04, 2020			U-1170
		6642245	Nov 04, 2020			U-248
		6642245	Nov 04, 2020			U-541
		6703396	Mar 09, 2021	DS DP		
		8592397	Jan 13, 2024	DP		U-1170
		8592397	Jan 13, 2024	DP		U-248
		8592397	Jan 13, 2024	DP		U-541
		8716264	Jan 13, 2024	DP		U-257
		9457036	Jan 13, 2024	DP		U-257
		9744181	Jan 13, 2024	DP		U-257

EMTRICITABINE; TENOFOVIR DISOPROXIL FUMARATE - TRUVADA

N 021752	002	6642245	Nov 04, 2020			U-1170
		6642245	Nov 04, 2020			U-248
		6642245	Nov 04, 2020			U-541
		6642245*PED	May 04, 2021			
		6703396	Mar 09, 2021	DS DP		
		6703396*PED	Sep 09, 2021			

EMTRICITABINE; TENOFOVIR DISOPROXIL FUMARATE - TRUVADA

N 021752	003	6642245	Nov 04, 2020			U-1170
		6642245	Nov 04, 2020			U-248
		6642245	Nov 04, 2020			U-541
		6642245*PED	May 04, 2021			
		6703396	Mar 09, 2021	DS DP		
		6703396*PED	Sep 09, 2021			

EMTRICITABINE; TENOFOVIR DISOPROXIL FUMARATE - TRUVADA

N 021752	004	6642245	Nov 04, 2020			U-1170
		6642245	Nov 04, 2020			U-248
		6642245	Nov 04, 2020			U-541
		6642245*PED	May 04, 2021			
		6703396	Mar 09, 2021	DS DP		
		6703396*PED	Sep 09, 2021			

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<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		
		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	NCE Aug 01, 2022
		10294215	Jan 07, 2033	DP	U-2087	ODE-151 Aug 01, 2024
		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	NCE Aug 01, 2022
		10294215	Jan 07, 2033	DP	U-2087	ODE-151 Aug 01, 2024
		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	NCE Jun 27, 2023
		8501758	Mar 04, 2031	DS DP		ODE-194 Jun 27, 2025
		8541575	Feb 26, 2030	DS DP	U-2335	
		8946250	Jul 23, 2029	DS DP		
		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	NCE Jun 27, 2023
		8501758	Mar 04, 2031	DS DP		ODE-194 Jun 27, 2025
		8541575	Feb 26, 2030	DS DP	U-2335	
		8946250	Jul 23, 2029	DS DP		
		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	

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<u>ENTRECTINIB - ROZLYTREK</u>						
N 212725 001	10231965	Feb 17, 2035	U-2617		NCE	Aug 15, 2024
	10231965	Feb 17, 2035	U-2618		ODE-265	Aug 15, 2026
	10398693	Jul 18, 2038	DP			
	10561651	Feb 19, 2035	U-2745			
	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 212725 002	10231965	Feb 17, 2035	U-2617		NCE	Aug 15, 2024
	10231965	Feb 17, 2035	U-2618		ODE-265	Aug 15, 2026
	10398693	Jul 18, 2038	DP			
	10561651	Feb 19, 2035	U-2745			
	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>ENZALUTAMIDE - XTANDI</u>						
N 203415 001	7709517	Aug 13, 2027	DS DP		I-786	Jul 13, 2021
	8183274	Aug 24, 2026	U-1281		I-808	Dec 16, 2022
	8183274	Aug 24, 2026	U-1588			
	8183274	Aug 24, 2026	U-2345			
	8183274	Aug 24, 2026	U-2708			
	9126941	May 15, 2026	U-1588			
	9126941	May 15, 2026	U-2345			
	9126941	May 15, 2026	U-2708			
<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 - TWINJECT 0.3</u>						
N 020800 001	7297136	Jan 18, 2025	DP			
	7621891	Feb 04, 2025	DP			
<u>EPINEPHRINE - TWINJECT 0.15</u>						
N 020800 002	7297136	Jan 18, 2025	DP			
	7621891	Feb 04, 2025	DP			
<u>EPINEPHRINE - ADRENALICK</u>						
N 020800 003	10166334	Jan 21, 2025	DP			
	7905352	Apr 12, 2027	DP			

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<u>EPINEPHRINE - ADRENALICK</u>						
N 020800	004	10166334	Jan 21, 2025	DP		
		7905352	Apr 12, 2027	DP		
<u>EPINEPHRINE - AUVI-O</u>						
N 201739	001	10314977	Nov 23, 2024	DP		
		10335549	Apr 30, 2025	DP		
		7731686	Jun 01, 2026	DP		
		7731690	Jan 15, 2025	DP		
		7749194	Oct 30, 2028	DP		
		7918823	Nov 23, 2024	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		
		8313466	Nov 23, 2024	DP		
		8361029	Nov 23, 2024	DP		
		8425462	Nov 23, 2024	DP		
		8608698	Nov 23, 2024	DP		
		8920377	Nov 23, 2024	DP		
		8926594	Mar 31, 2026	DP		
		9056170	Nov 23, 2024	DP		
		9149579	Jul 19, 2025		U-1758	
		9238108	Feb 20, 2027	DP		
		9259539	Feb 01, 2026	DP		
		9278182	Feb 01, 2026	DP		
		9724471	May 23, 2027	DP	U-2092	
		9737669	Nov 23, 2024	DP		
<u>EPINEPHRINE - AUVI-O</u>						
N 201739	002	10314977	Nov 23, 2024	DP		
		10335549	Apr 30, 2025	DP		
		7731686	Jun 01, 2026	DP		
		7731690	Jan 15, 2025	DP		
		7749194	Oct 30, 2028	DP		
		7918823	Nov 23, 2024	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		
		8313466	Nov 23, 2024	DP		
		8361029	Nov 23, 2024	DP		
		8425462	Nov 23, 2024	DP		
		8608698	Nov 23, 2024	DP		
		8920377	Nov 23, 2024	DP		
		8926594	Mar 31, 2026	DP		
		9056170	Nov 23, 2024	DP		
		9149579	Jul 19, 2025		U-1758	
		9238108	Feb 20, 2027	DP		
		9259539	Feb 01, 2026	DP		
		9278182	Feb 01, 2026	DP		
		9724471	May 23, 2027	DP	U-2092	
		9737669	Nov 23, 2024	DP		
<u>EPINEPHRINE - AUVI-O</u>						
N 201739	003	10314977	Nov 23, 2024	DP		
		10335549	Apr 30, 2025	DP		
		7731686	Jun 01, 2026	DP		
		7731690	Jan 15, 2025	DP		
		7749194	Oct 30, 2028	DP		
		7918823	Nov 23, 2024	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		

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<u>EPINEPHRINE - AUVI-Q</u>						
N 201739	003	8313466	Nov 23, 2024	DP		
		8361029	Nov 23, 2024	DP		
		8425462	Nov 23, 2024	DP		
		8608698	Nov 23, 2024	DP		
		8920377	Nov 23, 2024	DP		
		8926594	Mar 31, 2026	DP		
		9056170	Nov 23, 2024	DP		
		9149579	Jul 19, 2025		U-1758	
		9238108	Feb 20, 2027	DP		
		9259539	Feb 01, 2026	DP		
		9278182	Feb 01, 2026	DP		
		9724471	May 23, 2027	DP	U-2092	
		9737669	Nov 23, 2024	DP		
		9833573	Nov 23, 2024		U-2172	
<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	
		10039728	Aug 14, 2034		U-1828	
		9283197	Aug 15, 2034	DP	U-1828	
		9283197	Aug 15, 2034	DP	U-1829	
		9283197	Aug 15, 2034	DP	U-1830	
<u>EPINEPHRINE - PRIMATENE MIST</u>						
N 205920	001	8367734	Jan 26, 2026	DP		Nov 07, 2021
<u>EPINEPHRINE; LIDOCAINE HYDROCHLORIDE - LIDOSITE TOPICAL SYSTEM KIT</u>						
N 021504	001	6629968	Jun 30, 2020	DS DP		
		6635045	Jun 29, 2021	DS DP		
<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>ERAVACYCLINE DIHYDROCHLORIDE - XERAVA</u>						
N 211109	001	8796245	Aug 07, 2029		U-2380	Aug 27, 2023
		8906887	Dec 28, 2030	DP		Aug 27, 2028
<u>ERDAFITINIB - BALVERSA</u>						
N 212018	001	8895601	May 22, 2031	DS DP		Apr 12, 2024
		9464071	Apr 28, 2031		U-2518	
		9902714	Mar 26, 2035	DP		
<u>ERDAFITINIB - BALVERSA</u>						
N 212018	002	8895601	May 22, 2031	DS DP		Apr 12, 2024
		9464071	Apr 28, 2031		U-2518	
		9902714	Mar 26, 2035	DP		
<u>ERDAFITINIB - BALVERSA</u>						
N 212018	003	8895601	May 22, 2031	DS DP		Apr 12, 2024
		9464071	Apr 28, 2031		U-2518	
		9902714	Mar 26, 2035	DP		
<u>ERIBULIN MESYLATE - HALAVEN</u>						
N 201532	001	6214865	Jul 20, 2023	DS		Jan 28, 2023
		8097648	Jan 22, 2021		U-1096	
		RE46965	Jan 08, 2027	DP		

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<u>ERLOTINIB HYDROCHLORIDE - TARCEVA</u>						
N 021743 001	6900221	Nov 09, 2020	DS DP U-1046			
	6900221	Nov 09, 2020	DS DP U-1403			
	6900221	Nov 09, 2020	DS DP U-659			
	6900221	Nov 09, 2020	DS DP U-875			
	6900221*PED	May 09, 2021				
	7087613	Nov 09, 2020	U-1045			
	7087613	Nov 09, 2020	U-1403			
	7087613	Nov 09, 2020	U-659			
	7087613*PED	May 09, 2021				
<u>ERLOTINIB HYDROCHLORIDE - TARCEVA</u>						
N 021743 002	6900221	Nov 09, 2020	DS DP U-1046			
	6900221	Nov 09, 2020	DS DP U-1403			
	6900221	Nov 09, 2020	DS DP U-659			
	6900221	Nov 09, 2020	DS DP U-875			
	6900221*PED	May 09, 2021				
	7087613	Nov 09, 2020	U-1045			
	7087613	Nov 09, 2020	U-1403			
	7087613	Nov 09, 2020	U-659			
	7087613*PED	May 09, 2021				
<u>ERLOTINIB HYDROCHLORIDE - TARCEVA</u>						
N 021743 003	6900221	Nov 09, 2020	DS DP U-1046			
	6900221	Nov 09, 2020	DS DP U-1403			
	6900221	Nov 09, 2020	DS DP U-659			
	6900221	Nov 09, 2020	DS DP U-875			
	6900221*PED	May 09, 2021				
	7087613	Nov 09, 2020	U-1045			
	7087613	Nov 09, 2020	U-1403			
	7087613	Nov 09, 2020	U-659			
	7087613*PED	May 09, 2021				
<u>ERTUGLIFLOZIN - STEGLATRO</u>						
N 209803 001	8080580	Jul 13, 2030	DS DP U-2214		NCE	Dec 19, 2022
<u>ERTUGLIFLOZIN - STEGLATRO</u>						
N 209803 002	8080580	Jul 13, 2030	DS DP U-2214		NCE	Dec 19, 2022
<u>ERTUGLIFLOZIN; METFORMIN HYDROCHLORIDE - SEGLUOMET</u>						
N 209806 001	8080580	Jul 13, 2030	DS DP U-2214		NCE	Dec 19, 2022
	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		NCE	Dec 19, 2022
	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		NCE	Dec 19, 2022
	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		NCE	Dec 19, 2022
	9308204	Oct 21, 2030	DP			
	9439902	Oct 21, 2030	U-2214			
<u>ERTUGLIFLOZIN; SITAGLIPTIN PHOSPHATE - STEGLUJAN</u>						
N 209805 001	6699871	Jul 26, 2022	DS DP U-2214		NCE	Dec 19, 2022
	7326708	Nov 24, 2026	DS DP U-2214			
	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	6699871	Jul 26, 2022	DS DP U-2214		NCE	Dec 19, 2022
	7326708	Nov 24, 2026	DS DP U-2214			
	8080580	Jul 13, 2030	DS DP U-2214			
	9308204	Oct 21, 2030	DP			

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<u>ERTUGLIFLOZIN; SITAGLIPTIN PHOSPHATE - STEGLUJAN</u>						
N 209805 002	9439901	Oct 21, 2030	U-2214			
<u>ERYTHROMYCIN ETHYLSUCCINATE - ERYTHROMYCIN ETHYLSUCCINATE</u>						
A 211991 001					CGT	May 09, 2020
<u>ERYTHROMYCIN ETHYLSUCCINATE - ERYTHROMYCIN ETHYLSUCCINATE</u>						
A 211991 002					CGT	May 09, 2020
<u>ESCITALOPRAM OXALATE - LEXAPRO</u>						
N 021323 001	6916941	Aug 12, 2022	DS DP			
	7420069	Aug 12, 2022	DP			
<u>ESCITALOPRAM OXALATE - LEXAPRO</u>						
N 021323 002	6916941	Aug 12, 2022	DS DP			
	7420069	Aug 12, 2022	DP			
<u>ESCITALOPRAM OXALATE - LEXAPRO</u>						
N 021323 003	6916941	Aug 12, 2022	DS DP			
	7420069	Aug 12, 2022	DP			
<u>ESKETAMINE HYDROCHLORIDE - SPRAVATO</u>						
N 211243 001	8785500	Jul 09, 2031		U-2502	NCE*	Mar 05, 2024
	9592207	Mar 20, 2027		U-2502		
<u>ESLICARBAZEPINE ACETATE - APTIOM</u>						
N 022416 001	5753646	Jun 27, 2021	DS DP	U-2041		
	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	5753646	Jun 27, 2021	DS DP	U-2041		
	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	5753646	Jun 27, 2021	DS DP	U-2041		
	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	5753646	Jun 27, 2021	DS DP	U-2041		
	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 - BREVIBLOC IN PLASTIC CONTAINER</u>						
N 019386 004	6310094	Jan 12, 2021				
	6528540	Jan 12, 2021				

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<u>ESMOLOL HYDROCHLORIDE - BREVIBLOC DOUBLE STRENGTH IN PLASTIC CONTAINER</u>						
N 019386 005	6310094	Jan 12, 2021				
	6528540	Jan 12, 2021				
<u>ESMOLOL HYDROCHLORIDE - BREVIBLOC</u>						
N 019386 006	6310094	Jan 12, 2021				
	6528540	Jan 12, 2021				
<u>ESMOLOL HYDROCHLORIDE - BREVIBLOC</u>						
N 019386 007	6310094	Jan 12, 2021				
	6528540	Jan 12, 2021				
<u>ESMOLOL HYDROCHLORIDE - ESMOLOL HYDROCHLORIDE IN PLASTIC CONTAINER</u>						
N 205703 001	6310094	Jan 12, 2021	DP			
	6528540	Jan 12, 2021	DP			
	8829054	Mar 15, 2033	DP			
	8835505	Mar 15, 2033	DP			
<u>ESMOLOL HYDROCHLORIDE - ESMOLOL HYDROCHLORIDE DOUBLE STRENGTH IN PLASTIC CONTAINER</u>						
N 205703 002	6310094	Jan 12, 2021	DP			
	6528540	Jan 12, 2021	DP			
	8829054	Mar 15, 2033	DP			
	8835505	Mar 15, 2033	DP			
<u>ESOMEPRAZOLE MAGNESIUM - NEXIUM 24HR</u>						
N 204655 001	6428810*PED	May 03, 2020				
<u>ESOMEPRAZOLE MAGNESIUM - NEXIUM 24HR</u>						
N 207920 001	6428810*PED	May 03, 2020				
<u>ESOMEPRAZOLE MAGNESIUM; NAPROXEN - VIMOVO</u>						
N 022511 001	6926907	Feb 28, 2023	DP U-1052		NPP	Jul 06, 2020
	8557285	May 31, 2022	DP			
	8852636	May 31, 2022	DP U-1052			
	8858996	May 31, 2022	DP U-1052			
	8945621	Oct 17, 2031	U-1661			
	9161920	May 31, 2022	U-1760			
	9198888	May 31, 2022	U-1781			
	9220698	Mar 10, 2031	U-1781			
	9345695	May 31, 2022	DP			
	9393208	Sep 03, 2029	U-1781			
	9707181	May 31, 2022	DP			
<u>ESOMEPRAZOLE MAGNESIUM; NAPROXEN - VIMOVO</u>						
N 022511 002	6926907	Feb 28, 2023	DP U-1052		NPP	Jul 06, 2020
	8557285	May 31, 2022	DP			
	8852636	May 31, 2022	DP U-1052			
	8858996	May 31, 2022	DP U-1052			
	8945621	Oct 17, 2031	U-1661			
	9161920	May 31, 2022	U-1760			
	9198888	May 31, 2022	U-1781			
	9345695	May 31, 2022	DP			
	9393208	Sep 03, 2029	U-1781			
	9707181	May 31, 2022	DP			
<u>ESTRADIOL - VAGIFEM</u>						
N 020908 002	7018992	Sep 17, 2022	U-1023			
<u>ESTRADIOL - ELESTRIN</u>						
N 021813 001	7198801	Jun 25, 2022	DP			
	7470433	Aug 03, 2021	DP			
<u>ESTRADIOL - EVAMIST</u>						
N 022014 001	6978945	Jul 31, 2022	DP			
<u>ESTRADIOL - MINIVELLE</u>						
N 203752 001	6841716	Apr 27, 2020	DP			
	8231906	Jul 04, 2030	DS DP			
	9730900	Jul 10, 2028			U-2086	
	9833419	Jul 10, 2028	DP			

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<u>ESTRADIOL - MINIVELLE</u>						
N 203752	002	6841716	Apr 27, 2020	DP		
		8231906	Jul 04, 2030	DS DP		
		9730900	Jul 10, 2028		U-2086	
		9833419	Jul 10, 2028	DP		
<u>ESTRADIOL - MINIVELLE</u>						
N 203752	003	6841716	Apr 27, 2020	DP		
		8231906	Jul 04, 2030	DS DP		
		9730900	Jul 10, 2028		U-2086	
		9833419	Jul 10, 2028	DP		
<u>ESTRADIOL - MINIVELLE</u>						
N 203752	004	6841716	Apr 27, 2020	DP		
		8231906	Jul 04, 2030	DS DP		
		9730900	Jul 10, 2028		U-2086	
		9833419	Jul 10, 2028	DP		
<u>ESTRADIOL - MINIVELLE</u>						
N 203752	005	6841716	Apr 27, 2020	DP		
		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	Dec 20, 2033		U-2316	NP May 29, 2021
		10258630	Dec 20, 2033		U-2317	
		10398708	Dec 20, 2033		U-2317	
		10398708	Dec 20, 2033		U-2614	
		10471072	Jun 18, 2033		U-2316	
		10471072	Jun 18, 2033		U-2317	
		10537581	Nov 21, 2032	DP	U-2316	
		10537581	Nov 21, 2032	DP	U-2317	
		9180091	Dec 20, 2033	DP	U-2316	
		9180091	Dec 20, 2033	DP	U-2317	
		9289382	Nov 21, 2032	DP		
<u>ESTRADIOL - IMVEXXY</u>						
N 208564	002	10258630	Dec 20, 2033		U-2316	NP May 29, 2021
		10258630	Dec 20, 2033		U-2317	
		10398708	Dec 20, 2033		U-2317	
		10398708	Dec 20, 2033		U-2614	
		10471072	Jun 18, 2033		U-2316	
		10471072	Jun 18, 2033		U-2317	
		10537581	Nov 21, 2032	DP	U-2316	
		10537581	Nov 21, 2032	DP	U-2317	
		9180091	Dec 20, 2033	DP	U-2316	
		9180091	Dec 20, 2033	DP	U-2317	
		9289382	Nov 21, 2032	DP		
<u>ESTRADIOL ACETATE - FEMTRACE</u>						
N 021633	001	6962908	Dec 21, 2021	DP		
		7572779	Oct 02, 2025		U-904	
		7799771	Dec 21, 2021	DP		
<u>ESTRADIOL ACETATE - FEMTRACE</u>						
N 021633	002	6962908	Dec 21, 2021	DP		
		7572779	Oct 02, 2025		U-904	
		7799771	Dec 21, 2021	DP		
<u>ESTRADIOL ACETATE - FEMTRACE</u>						
N 021633	003	6962908	Dec 21, 2021	DP		
		7572779	Oct 02, 2025		U-904	
		7799771	Dec 21, 2021	DP		
<u>ESTRADIOL; ESTRADIOL; NORGESTIMATE - PREFEST</u>						
N 021040	001	6747019	Mar 20, 2020		U-311	
		7320970	Mar 30, 2020	DP	U-844	

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<u>ESTRADIOL; PROGESTERONE - BIJUVA</u>						
N 210132	001	10052386	Nov 21, 2032	DP	NP	Oct 28, 2021
		10206932	Nov 21, 2032	U-2439		
		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>ESTROGENS, CONJUGATED SYNTHETIC B - ENJUVA</u>						
N 021443	001	6660726	Mar 08, 2021	DS DP U-904		
		6660726	Mar 08, 2021	DS DP U-905		
		6855703	Feb 12, 2021	DS DP U-904		
		6855703	Feb 12, 2021	DS DP U-905		
<u>ESTROGENS, CONJUGATED SYNTHETIC B - ENJUVA</u>						
N 021443	002	6660726	Mar 08, 2021	DS DP U-904		
		6660726	Mar 08, 2021	DS DP U-905		
		6855703	Feb 12, 2021	DS DP U-904		
		6855703	Feb 12, 2021	DS DP U-905		
<u>ESTROGENS, CONJUGATED SYNTHETIC B - ENJUVA</u>						
N 021443	003	6660726	Mar 08, 2021	DS DP U-904		
		6660726	Mar 08, 2021	DS DP U-905		
		6855703	Feb 12, 2021	DS DP U-904		
		6855703	Feb 12, 2021	DS DP U-905		
<u>ESTROGENS, CONJUGATED SYNTHETIC B - ENJUVA</u>						
N 021443	004	6660726	Mar 08, 2021	DS DP U-904		
		6660726	Mar 08, 2021	DS DP U-905		
		6855703	Feb 12, 2021	DS DP U-904		
		6855703	Feb 12, 2021	DS DP U-905		
<u>ESTROGENS, CONJUGATED SYNTHETIC B - ENJUVA</u>						
N 021443	005	6660726	Mar 08, 2021	DS DP U-904		
		6660726	Mar 08, 2021	DS DP U-905		
		6855703	Feb 12, 2021	DS DP U-904		
		6855703	Feb 12, 2021	DS DP U-905		
<u>ETELCALCETIDE - PARSABIV</u>						
N 208325	001	10344765	Jun 27, 2034	DP	NCE	Feb 07, 2022
		8377880	Jul 29, 2030	DS DP		
		8999932	Jul 29, 2030	DS DP U-2014		
		9278995	Jul 29, 2030	DS		
		9701712	Jul 29, 2030	DS DP U-2014		
		9820938	Jun 27, 2034	DP		
<u>ETELCALCETIDE - PARSABIV</u>						
N 208325	002	10344765	Jun 27, 2034	DP	NCE	Feb 07, 2022
		8377880	Jul 29, 2030	DS DP		
		8999932	Jul 29, 2030	DS DP U-2014		
		9278995	Jul 29, 2030	DS		
		9701712	Jul 29, 2030	DS DP U-2014		
		9820938	Jun 27, 2034	DP		
<u>ETELCALCETIDE - PARSABIV</u>						
N 208325	003	10344765	Jun 27, 2034	DP	NCE	Feb 07, 2022
		8377880	Jul 29, 2030	DS DP		
		8999932	Jul 29, 2030	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	NCE	Sep 19, 2021
		10364431	Mar 14, 2034	U-1918	ODE-122	Sep 19, 2023

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<u>ETEPLIRSEN - EXONDYS 51</u>						
N 206488 001	10364431	Mar 14, 2034	U-1919			
	10533174	May 04, 2021	DP			
	8486907	Jun 28, 2025	U-1904	Y		
	9018368	Jun 28, 2025	DS DP			
	9243245	Oct 27, 2028	DS U-2097			
	9243245	Oct 27, 2028	DS U-2098			
	9416361	May 04, 2021	DS			
	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	Jun 28, 2025	DP			
<u>ETEPLIRSEN - EXONDYS 51</u>						
N 206488 002	10337003	Mar 14, 2034	U-1918		NCE	Sep 19, 2021
	10364431	Mar 14, 2034	U-1918		ODE-122	Sep 19, 2023
	10364431	Mar 14, 2034	U-1919			
	10533174	May 04, 2021	DP			
	8486907	Jun 28, 2025	U-1904	Y		
	9018368	Jun 28, 2025	DS DP			
	9243245	Oct 27, 2028	DS U-2097			
	9243245	Oct 27, 2028	DS U-2098			
	9416361	May 04, 2021	DS			
	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	Jun 28, 2025	DP			
<u>ETHINYL ESTRADIOL; ETHINYL ESTRADIOL; LEVONORGESTREL - SEASONIQUE</u>						
N 021840 001	7320969	Jan 30, 2024	U-828			
	7615545	Jun 15, 2023	U-1			
	7855190	Dec 05, 2028	U-1			
	7858605	Jun 23, 2023	DP			
<u>ETHINYL ESTRADIOL; LEVONORGESTREL - PREVEN EMERGENCY CONTRACEPTIVE KIT</u>						
N 020946 001	6156742	Dec 05, 2020	U-374			
<u>ETHINYL ESTRADIOL; LEVONORGESTREL - LOSEASONIQUE</u>						
N 022262 001	7615545	Jun 15, 2023	U-1			
	7855190	Dec 05, 2028	U-1			
	7858605	Jun 23, 2023	DP			
<u>ETHINYL ESTRADIOL; LEVONORGESTREL - TWIRLA</u>						
N 204017 001	7045145	Mar 14, 2021	DP U-2751		NP	Feb 14, 2023
	7384650	Mar 14, 2021	DP			
	8221784	Mar 14, 2021	DP			
	8221785	Mar 14, 2021	DP			
	8246978	Aug 26, 2028	DP			
	8747888	Jul 10, 2028	DP			
	8883196	Nov 22, 2020	DP U-2751			
	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	6716814	Aug 16, 2021	DS DP			
<u>ETHINYL ESTRADIOL; NORETHINDRONE ACETATE - LO LOESTRIN FE</u>						
N 022501 001	7704984	Feb 02, 2029	U-1090			
<u>ETHINYL ESTRADIOL; NORETHINDRONE ACETATE - TAYTULLA</u>						
N 204426 001	6652880	Mar 29, 2020	DP			

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<u>ETHINYL ESTRADIOL; NORETHINDRONE ACETATE - LO MINASTRIN FE</u>						
N 204654	001	7704984	Feb 02, 2029	U-1		
<u>ETHINYL ESTRADIOL; SEGESTERONE ACETATE - ANNOVERA</u>						
N 209627	001				NCE	Aug 10, 2023
<u>ETHIODIZED OIL - LIPIODOL</u>						
N 009190	001				ODE-64	Apr 04, 2021
<u>ETONOGESTREL - IMPLANON</u>						
N 021529	001	9757552	Jul 28, 2030	DP U-1		
<u>ETONOGESTREL - NEXPLANON</u>						
N 021529	002	8722037	Sep 28, 2027	DP		
		8888745	Aug 28, 2026	DP		
		9757552	Jul 28, 2030	DP U-1		
<u>ETRAVIRINE - INTELENCE</u>						
N 022187	001	6878717*PED	May 05, 2020		NPP	Jul 16, 2021
		7037917	Dec 13, 2020	DS DP U-1016	PED	Jan 16, 2022
		7037917	Dec 13, 2020	DS DP U-1237		
		7037917	Dec 13, 2020	DS DP U-2354		
		7037917	Dec 13, 2020	DS DP U-256		
		7037917*PED	Jun 13, 2021			
		8003789*PED	May 01, 2020			
<u>ETRAVIRINE - INTELENCE</u>						
N 022187	002	6878717*PED	May 05, 2020		NPP	Jul 16, 2021
		7037917	Dec 13, 2020	DS DP U-1016	PED	Jan 16, 2022
		7037917	Dec 13, 2020	DS DP U-1237		
		7037917	Dec 13, 2020	DS DP U-2354		
		7037917	Dec 13, 2020	DS DP U-256		
		7037917*PED	Jun 13, 2021			
		8003789*PED	May 01, 2020			
<u>ETRAVIRINE - INTELENCE</u>						
N 022187	003	6878717*PED	May 05, 2020		NPP	Jul 16, 2021
		7037917	Dec 13, 2020	DS DP U-1237	PED	Jan 16, 2022
		7037917	Dec 13, 2020	DS DP U-2354		
		7037917*PED	Jun 13, 2021			
		8003789*PED	May 01, 2020			
<u>EVEROLIMUS - ZORTRESS</u>						
N 021560	001	5665772*PED	Mar 09, 2020			
<u>EVEROLIMUS - ZORTRESS</u>						
N 021560	002	5665772*PED	Mar 09, 2020			
<u>EVEROLIMUS - ZORTRESS</u>						
N 021560	003	5665772*PED	Mar 09, 2020			
<u>EVEROLIMUS - AFINITOR</u>						
N 022334	001	8410131	Nov 01, 2025	U-1368	ODE-108	Feb 26, 2023
		8410131*PED	May 01, 2026			
		8436010	Feb 22, 2022	U-1396		
		8436010*PED	Aug 22, 2022			
		8778962	Feb 18, 2022	U-1541		
		8778962*PED	Aug 18, 2022			
		9006224	Jul 01, 2028	U-1681		
<u>EVEROLIMUS - AFINITOR</u>						
N 022334	002	8410131	Nov 01, 2025	U-1368	ODE-108	Feb 26, 2023
		8410131*PED	May 01, 2026			
		8436010	Feb 22, 2022	U-1396		
		8436010*PED	Aug 22, 2022			
		8778962	Feb 18, 2022	U-1541		
		8778962*PED	Aug 18, 2022			
		9006224	Jul 01, 2028	U-1681		

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<u>EVEROLIMUS - AFINITOR</u>						
N 022334 003	8410131	Nov 01, 2025	U-1368		ODE-108	Feb 26, 2023
	8410131*PED	May 01, 2026				
	8436010	Feb 22, 2022	U-1396			
	8436010*PED	Aug 22, 2022				
	8778962	Feb 18, 2022	U-1541			
	8778962*PED	Aug 18, 2022				
	9006224	Jul 01, 2028	U-1681			
<u>EVEROLIMUS - AFINITOR</u>						
N 022334 004	8410131	Nov 01, 2025	U-1368		ODE-108	Feb 26, 2023
	8410131*PED	May 01, 2026				
	8436010	Feb 22, 2022	U-1396			
	8436010*PED	Aug 22, 2022				
	8778962	Feb 18, 2022	U-1541			
	8778962*PED	Aug 18, 2022				
	9006224	Jul 01, 2028	U-1681			
<u>EVEROLIMUS - AFINITOR DISPERZ</u>						
N 203985 001	8617598	Sep 27, 2022	DP		I-773	Apr 10, 2021
	8617598*PED	Mar 27, 2023			ODE-169	Apr 10, 2025
	8778962	Feb 18, 2022	U-1541			
	8778962	Feb 18, 2022	U-2280			
	8778962*PED	Aug 18, 2022				
<u>EVEROLIMUS - AFINITOR DISPERZ</u>						
N 203985 002	8617598	Sep 27, 2022	DP		I-773	Apr 10, 2021
	8617598*PED	Mar 27, 2023			ODE-169	Apr 10, 2025
	8778962	Feb 18, 2022	U-1541			
	8778962	Feb 18, 2022	U-2280			
	8778962*PED	Aug 18, 2022				
<u>EVEROLIMUS - AFINITOR DISPERZ</u>						
N 203985 003	8617598	Sep 27, 2022	DP		I-773	Apr 10, 2021
	8617598*PED	Mar 27, 2023			ODE-169	Apr 10, 2025
	8778962	Feb 18, 2022	U-1541			
	8778962	Feb 18, 2022	U-2280			
	8778962*PED	Aug 18, 2022				
<u>EXENATIDE - BYDUREON BCISE</u>						
N 209210 001	6414126	Oct 04, 2020	U-2588		NP	Oct 20, 2020
	6479065	Aug 10, 2020	DP			
	6515117	Oct 04, 2020	U-2588			
	6667061	May 25, 2020	DP			
	6824822	Oct 09, 2022	DP			
	6872700	Jan 14, 2020	U-2589			
	6872700	Jan 14, 2020	U-2590			
	6872700	Jan 14, 2020	U-2591			
	6936590	Oct 04, 2020	U-2588			
	7223440	Aug 31, 2021	DP			
	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-2592			
	7563871	Apr 15, 2024	DP			
	7612176	Apr 13, 2025	DP U-2589			
	7612176	Apr 13, 2025	DP U-2590			
	7612176	Apr 13, 2025	DP U-2592			
	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			
	8361972	Mar 21, 2028	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-2598			
	8461105	Apr 13, 2025	DP U-2589			

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<u>EXENATIDE - BYDUREON BCISE</u>						
N 209210 001	8461105	Apr 13, 2025	DP U-2590			
	8461105	Apr 13, 2025	DP U-2597			
	8461105	Apr 13, 2025	DP U-2598			
	8501698	Jun 20, 2027	U-2588			
	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-2600			
	8895033	Oct 04, 2030	DP U-2601			
	8895033	Oct 04, 2030	DP U-2602			
	8906851	Aug 18, 2026	U-2589			
	8906851	Aug 18, 2026	U-2590			
	8906851	Aug 18, 2026	U-2593			
	8906851	Aug 18, 2026	U-2597			
	9198925	Oct 04, 2020	U-2588			
	9238076	Apr 15, 2024	DP U-2589			
	9238076	Apr 15, 2024	DP U-2590			
	9238076	Apr 15, 2024	DP U-2597			
	9238076	Apr 15, 2024	DP U-2599			
	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			
<u>EXENATIDE SYNTHETIC - BYETTA</u>						
N 021773 001	6872700	Jan 14, 2020	U-654			
	6902744	Jan 14, 2020	DP			
<u>EXENATIDE SYNTHETIC - BYETTA</u>						
N 021773 002	6872700	Jan 14, 2020	U-654			
	6902744	Jan 14, 2020	DP			
<u>EXENATIDE SYNTHETIC - BYDUREON</u>						
N 022200 001	6414126	Oct 04, 2020	DS DP U-2588		M-212	Oct 20, 2020
	6479065	Aug 10, 2020	DP		M-224	Apr 02, 2021
	6495164	May 25, 2020	DP			
	6515117	Oct 04, 2020	DS DP U-2588			
	6667061	May 25, 2020	DP			
	6824822	Oct 09, 2022	DP			
	6872700	Jan 14, 2020	U-2588			
	6872700	Jan 14, 2020	U-2589			
	6872700	Jan 14, 2020	U-2590			
	6872700	Jan 14, 2020	U-2591			
	6936590	Oct 04, 2020	U-2588			
	7223440	Aug 31, 2021	DP			
	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-2592			
	7563871	Apr 15, 2024	DP			
	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-2592			
	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			
	8361972	Mar 21, 2028	U-2588			
	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-2598			
	8461105	Apr 13, 2025	DP U-2588			

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<u>EXENATIDE SYNTHETIC - BYDUREON</u>						
N 022200 001	8461105	Apr 13, 2025	DP U-2589			
	8461105	Apr 13, 2025	DP U-2590			
	8461105	Apr 13, 2025	DP U-2598			
	8501698	Jun 20, 2027	DP U-2588			
	8906851	Aug 18, 2026	U-2588			
	8906851	Aug 18, 2026	U-2589			
	8906851	Aug 18, 2026	U-2590			
	8906851	Aug 18, 2026	U-2593			
	9198925	Oct 04, 2020	U-2588			
	9238076	Apr 15, 2024	DP U-2588			
	9238076	Apr 15, 2024	DP U-2589			
	9238076	Apr 15, 2024	DP U-2590			
	9238076	Apr 15, 2024	DP U-2599			
	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			
<u>EXENATIDE SYNTHETIC - BYDUREON PEN</u>						
N 022200 002	6414126	Oct 04, 2020	DS DP U-2588		M-224	Apr 02, 2021
	6479065	Aug 10, 2020	DP			
	6495164	May 25, 2020	DP			
	6515117	Oct 04, 2020	DS DP U-2588			
	6667061	May 25, 2020	DP			
	6824822	Oct 09, 2022	DP			
	6872700	Jan 14, 2020	U-2588			
	6872700	Jan 14, 2020	U-2589			
	6872700	Jan 14, 2020	U-2590			
	6872700	Jan 14, 2020	U-2591			
	6936590	Oct 04, 2020	U-2588			
	7223440	Aug 31, 2021	DP			
	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-2592			
	7563871	Apr 15, 2024	DP			
	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-2592			
	8216180	Jan 12, 2028	DP			
	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			
	8361972	Mar 21, 2028	U-2588			
	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-2598			
	8439864	Mar 25, 2028	DP			
	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-2598			
	8501698	Jun 20, 2027	DP U-2588			
	8690837	May 19, 2029	DP			
	8721615	Jan 18, 2030	DP			
	8758292	Nov 12, 2027	DP			
	8827963	Feb 04, 2029	DP			
	8906851	Aug 18, 2026	U-2588			
	8906851	Aug 18, 2026	U-2589			
	8906851	Aug 18, 2026	U-2590			

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<u>EXENATIDE SYNTHETIC - BYDUREON PEN</u>						
N 022200 002	8906851	Aug 18, 2026	U-2593			
	8998876	Jan 07, 2030	DP			
	9198925	Oct 04, 2020	U-2588			
	9238076	Apr 15, 2024	DP U-2588			
	9238076	Apr 15, 2024	DP U-2589			
	9238076	Apr 15, 2024	DP U-2590			
	9238076	Apr 15, 2024	DP U-2599			
	9320853	Mar 25, 2028	DP			
	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			
<u>EZETIMIBE - ZETIA</u>						
N 021445 001	7030106	Jan 25, 2022	DP			
	7612058	Oct 30, 2025	U-1027			
	7612058	Oct 30, 2025	U-1173			
	7612058*PED	Apr 30, 2026				
<u>FAMOTIDINE - PEPCID AC</u>						
N 020801 002	6814978	Aug 26, 2021	DP			
<u>FAMOTIDINE; IBUPROFEN - DUEXIS</u>						
N 022519 001	8067033	Jul 18, 2026	DP			
	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			
<u>FEBUXOSTAT - ULORIC</u>						
N 021856 001	7361676	Mar 08, 2024	DP		M-205	Aug 15, 2020
	8372872	Sep 08, 2031	U-1346			
	9107912	Sep 08, 2031	U-1346			
<u>FEBUXOSTAT - ULORIC</u>						
N 021856 002	7361676	Mar 08, 2024	DP		M-205	Aug 15, 2020
	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		NCE	Aug 16, 2024
	7528143	Dec 16, 2026	DS		ODE-259	Aug 16, 2026
	7825246	Dec 16, 2026	DS			
	8138199	Jun 30, 2028	U-2607			
<u>FENOFIBRATE - TRIGLIDE</u>						
N 021350 001	6696084	Sep 11, 2021	DS DP U-680			
<u>FENOFIBRATE - TRIGLIDE</u>						
N 021350 002	6696084	Sep 11, 2021	DS DP U-680			
<u>FENOFIBRATE - TRICOR</u>						
N 021656 001	6375986	Sep 21, 2020	DP U-615			
	7276249	Feb 21, 2023	DP			
	7320802	Feb 21, 2023	U-847			
<u>FENOFIBRATE - TRICOR</u>						
N 021656 002	6375986	Sep 21, 2020	DP U-615			
	7276249	Feb 21, 2023	DP			
	7320802	Feb 21, 2023	U-847			
<u>FENOFIBRATE - ANTARA (MICRONIZED)</u>						
N 021695 001	7101574	Aug 20, 2020	DS DP			
	7863331	Aug 08, 2020	U-1106			
	7863331	Aug 08, 2020	U-1107			

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<u>FENOFIBRATE - ANTARA (MICRONIZED)</u>						
N 021695	003	7101574	Aug 20, 2020	DS DP		
		7863331	Aug 08, 2020		U-1106	
		7863331	Aug 08, 2020		U-1107	
<u>FENOFIBRATE - ANTARA (MICRONIZED)</u>						
N 021695	004	8026281	Apr 22, 2025		U-1447	
		8026281	Apr 22, 2025		U-1448	
<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>FENOFIBRATE - FENOGLIDE</u>						
N 022118	001	7658944	Dec 09, 2024	DP		
		8124125	Oct 01, 2024	DP	U-1234	
		8481078	Oct 01, 2024	DP	U-1416	
		9173847	Oct 01, 2024	DP		
<u>FENOFIBRATE - FENOGLIDE</u>						
N 022118	002	7658944	Dec 09, 2024	DP		
		8124125	Oct 01, 2024	DP	U-1234	
		8481078	Oct 01, 2024	DP	U-1416	
		9173847	Oct 01, 2024	DP		
<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		
		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		
		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		
		8486972	Apr 27, 2030	DP		
		8486973	Apr 27, 2030		U-55	
		8835459	Jan 25, 2027	DP		

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<u>FENTANYL - SUBSYS</u>						
N 202788 003	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 004	10016403	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			
	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			
	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			
	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 CITRATE - FENTORA</u>						
N 021947 001	7862832	Jun 15, 2028	DP			
	7862833	Jun 15, 2028	DP			
	8092832	Dec 30, 2024	DP			
<u>FENTANYL CITRATE - FENTORA</u>						
N 021947 002	7862832	Jun 15, 2028	DP			
	7862833	Jun 15, 2028	DP			
	8092832	Dec 30, 2024	DP			
	8119158	Dec 30, 2024	DP			
<u>FENTANYL CITRATE - FENTORA</u>						
N 021947 003	7862832	Jun 15, 2028	DP			
	7862833	Jun 15, 2028	DP			
	8092832	Dec 30, 2024	DP			
	8119158	Dec 30, 2024	DP			
<u>FENTANYL CITRATE - FENTORA</u>						
N 021947 004	7862832	Jun 15, 2028	DP			
	7862833	Jun 15, 2028	DP			
	8092832	Dec 30, 2024	DP			

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<u>FENTANYL CITRATE - FENTORA</u>						
N 021947	004	8119158	Dec 30, 2024	DP		
<u>FENTANYL CITRATE - FENTORA</u>						
N 021947	005	7862832	Jun 15, 2028	DP		
		7862833	Jun 15, 2028	DP		
		8092832	Dec 30, 2024	DP		
		8119158	Dec 30, 2024	DP		
<u>FENTANYL CITRATE - ONSOLIS</u>						
N 022266	001	7579019	Jan 22, 2020		U-767	
		9597288	Jul 23, 2027	DP	U-767	
<u>FENTANYL CITRATE - ONSOLIS</u>						
N 022266	002	7579019	Jan 22, 2020		U-767	
		9597288	Jul 23, 2027	DP	U-767	
<u>FENTANYL CITRATE - ONSOLIS</u>						
N 022266	003	7579019	Jan 22, 2020		U-767	
		9597288	Jul 23, 2027	DP	U-767	
<u>FENTANYL CITRATE - ONSOLIS</u>						
N 022266	004	7579019	Jan 22, 2020		U-767	
		9597288	Jul 23, 2027	DP	U-767	
<u>FENTANYL CITRATE - ONSOLIS</u>						
N 022266	005	7579019	Jan 22, 2020		U-767	
		9597288	Jul 23, 2027	DP	U-767	
<u>FENTANYL CITRATE - LAZANDA</u>						
N 022569	001	8216604	Oct 03, 2024		U-767	
		8889176	Jan 16, 2024		U-767	
		9078814	Jan 08, 2024	DP		
		9731869	Jan 26, 2032	DP		
		9814705	Jan 08, 2024	DP		
<u>FENTANYL CITRATE - LAZANDA</u>						
N 022569	002	8216604	Oct 03, 2024		U-767	
		8889176	Jan 16, 2024		U-767	
		9078814	Jan 08, 2024	DP		
		9731869	Jan 26, 2032	DP		
		9814705	Jan 08, 2024	DP		
<u>FENTANYL CITRATE - LAZANDA</u>						
N 022569	003	9731869	Jan 26, 2032	DP		
		9814705	Jan 08, 2024	DP		
<u>FENTANYL HYDROCHLORIDE - IONSYS</u>						
N 021338	001	6881208	Apr 19, 2022		U-736	
		6975902	Apr 01, 2024	DP		
		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	10519252	Oct 20, 2023	DS	U-2709	
		10519252	Oct 20, 2023	DS	U-2710	
		10519252	Oct 20, 2023	DS	U-2711	
		10519252	Oct 20, 2023	DS	U-2712	
		7612109	Feb 05, 2024	DS DP		
		7754702	Feb 15, 2028	DP	U-1432	
		7754702	Feb 15, 2028	DP	U-2555	
		7754702	Feb 15, 2028	DP	U-2556	
		7754702	Feb 15, 2028	DP	U-2557	
		8895612	Jan 08, 2027	DP	U-1620	
		9376505	Oct 20, 2023	DS DP		

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<u>FERRIC CITRATE - AURYXIA</u>						
N 205874 001	10300039	Jul 21, 2030		U-2549	I-790	Nov 06, 2020
	5753706	Feb 03, 2021		DP U-1577		
	7767851	Feb 18, 2024	DS DP			
	8093423	Apr 21, 2026		U-1577		
	8299298	Feb 18, 2024		DP		
	8338642	Feb 18, 2024	DS DP	U-1577		
	8609896	Feb 18, 2024		DP		
	8754257	Feb 18, 2024		DP		
	8754258	Feb 18, 2024		DP		
	8846976	Feb 18, 2024		U-1577		
	8901349	Feb 18, 2024		U-1577		
	9050316	Feb 18, 2024		U-1577		
	9328133	Feb 18, 2024	DS DP	U-1577		
	9387191	Jul 21, 2030		DP		
	9757416	Feb 18, 2024	DS DP	U-1577		
<u>FERRIC DERISOMALTOSE - MONOFERRIC</u>						
N 208171 001	10414831	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	NCE	Jul 25, 2024
	9248148	Mar 29, 2031		U-2603		
	9802973	Oct 23, 2035	DS DP	U-2603		
<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		
<u>FERUMOXYTOL - FERAHEME</u>						
N 022180 001	6599498	Jun 30, 2023	DS DP		I-767	Feb 02, 2021
	7553479	Mar 08, 2020	DS DP			
	7871597	Mar 08, 2020	DS DP			
	8501158	Mar 08, 2020		U-1422		
	8591864	Mar 08, 2020		DP		
	8926947	Mar 08, 2020	DS DP			
<u>FESOTERODINE FUMARATE - TOVIAZ</u>						
N 022030 001	6858650	Jul 03, 2022	DS	U-913		
	7807715	Jun 07, 2027		DP U-913		
	8088398	Jun 07, 2027		DP U-913		
	8501723	Jun 07, 2027		DP		
<u>FESOTERODINE FUMARATE - TOVIAZ</u>						
N 022030 002	6858650	Jul 03, 2022	DS	U-913		
	7807715	Jun 07, 2027		DP U-913		
	8088398	Jun 07, 2027		DP U-913		
	8501723	Jun 07, 2027		DP		
<u>FEXOFENADINE HYDROCHLORIDE - CHILDREN'S ALLEGRA ALLERGY</u>						
N 021909 002	6723348	Nov 26, 2021		DP U-1466		
<u>FEXOFENADINE HYDROCHLORIDE - CHILDREN'S ALLEGRA HIVES</u>						
N 021909 003	6723348	Nov 26, 2021		DP		
<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>FEXOFENADINE HYDROCHLORIDE; PSEUDOEPHEDRINE HYDROCHLORIDE - ALLEGRA-D 24 HOUR ALLERGY AND CONGESTION</u>						
N 021704 002	6613357	Dec 25, 2020		DP U-1159		
<u>FIDAXOMICIN - DIFICID</u>						
N 201699 001	7378508	Jul 31, 2027	DS DP		NPP	Jan 24, 2023
	7863249	Jul 31, 2027	DS DP		PED	Jul 24, 2023
	7906489	Mar 04, 2027		U-2741		

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<u>FIDAXOMICIN - DIFICID</u>						
N 201699 001	7906489	Mar 04, 2027	U-319			
	7906489*PED	Sep 04, 2027				
	8586551	Jul 15, 2023	DS DP			
	8586551*PED	Jan 15, 2024				
	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		NP	Jan 24, 2023
	7378508*PED	Jan 31, 2028			PED	Jul 24, 2023
	7863249	Jul 31, 2027	DP			
	7863249*PED	Jan 31, 2028				
	7906489	Mar 04, 2027	U-2741			
	7906489*PED	Sep 04, 2027				
	8586551	Jul 23, 2023	DS DP			
	8586551*PED	Jan 23, 2024				
	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			
	9504691	Nov 21, 2033	DP U-1679			
<u>FINGOLIMOD HYDROCHLORIDE - GILENYA</u>						
N 022527 001	10543179	Dec 25, 2027	U-2719		NPP	May 11, 2021
	8324283	Mar 29, 2026	DP		PED	Nov 11, 2021
	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		NS	May 11, 2021
	9592208*PED	Sep 30, 2032			PED	Nov 11, 2021
<u>FISH OIL TRIGLYCERIDES - OMEGAVEN</u>						
N 210589 001	10350186	Nov 05, 2024	U-2585		NCE	Jul 27, 2023
	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	10350186	Nov 05, 2024	U-2585		NCE	Jul 27, 2023
	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					NCE	Jul 13, 2021
<u>FISH OIL; MEDIUM CHAIN TRIGLYCERIDES; OLIVE OIL; SOYBEAN OIL - SMOFLIPID 20%</u>						
N 207648 002					NCE	Jul 13, 2021
<u>FISH OIL; MEDIUM CHAIN TRIGLYCERIDES; OLIVE OIL; SOYBEAN OIL - SMOFLIPID 20%</u>						
N 207648 003					NCE	Jul 13, 2021
<u>FISH OIL; MEDIUM CHAIN TRIGLYCERIDES; OLIVE OIL; SOYBEAN OIL - SMOFLIPID 20%</u>						
N 207648 004					NCE	Jul 13, 2021
<u>FLIBANSERIN - ADDYI</u>						
N 022526 001	7151103	May 09, 2023	U-1734		NCE	Aug 18, 2020
	7420057	Aug 01, 2022	DS DP			
	8227471	May 09, 2023	U-1734			
	9468639	Oct 16, 2022	U-1734			
<u>FLORBETABEN F-18 - NEURACEQ</u>						
N 204677 001	7807135	Mar 18, 2029	DS DP U-1497			

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<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>FLUCICLOVINE F-18 - AXUMIN</u>						
N 208054 001	10010632	Nov 28, 2026	DP		NCE	May 27, 2021
	10124079	Dec 30, 2035	U-2450			
	5808146	Nov 09, 2020	DS			
	9387266	Nov 28, 2026	U-1879			
<u>FLUDARABINE PHOSPHATE - OFORTA</u>						
N 022273 001	7148207	Dec 20, 2022	DP U-944			
<u>FLUOCINOLONE ACETONIDE - ILLUVIEN</u>						
N 201923 001	6375972	Apr 26, 2020	DP U-1597			
	8871241	Aug 12, 2027	DP			
<u>FLUOCINOLONE ACETONIDE - YUTIQ</u>						
N 210331 001	6375972	Apr 26, 2020	DP U-708		NP	Nov 12, 2021
	8574613	Apr 26, 2020	DP			
	8574659	Apr 26, 2020	DP			
	8871241	Aug 12, 2027	DP			
	9192579	Apr 26, 2020	DP			
	9849085	Apr 26, 2020	U-708			
<u>FLUOCINOLONE ACETONIDE; HYDROQUINONE; TRETINOIN - TRI-LUMA</u>						
N 021112 001	7915243	Sep 08, 2023	DP			
	7939516	Sep 08, 2023	DP			
	8247395	Oct 25, 2022	DP			
	8653053	Oct 25, 2022	DP			
<u>FLUOCINONIDE - VANOS</u>						
N 021758 001	6765001	Dec 21, 2021	DP			
	7220424	Jan 07, 2023	U-861			
	7794738	Sep 11, 2022	U-1084			
	8232264	Mar 09, 2023	DP			
<u>FLUORODOPA F-18 - FLUORODOPA F18</u>						
N 200655 001					NCE W	Oct 10, 2024 Oct 10, 2024
<u>FLUOROURACIL - CARAC</u>						
N 020985 001	6670335	Jun 02, 2021	DP U-68			
<u>FLUOROURACIL - TOLAK</u>						
N 022259 001	7169401	Jul 18, 2023	DP			
<u>FLUTEMETAMOL F-18 - VIZAMYL</u>						
N 203137 001	7270800	Sep 03, 2025	DS DP U-336			
	7351401	Jan 24, 2023	DS DP U-336			
	8236282	May 21, 2024	DS DP			
	8691185	Jan 24, 2023	U-336			
	8916131	Sep 16, 2028	DP			
<u>FLUTEMETAMOL F-18 - VIZAMYL</u>						
N 203137 002	7270800	Sep 03, 2025	DS DP U-336			
	7351401	Jan 24, 2023	DS DP U-336			
	8236282	May 21, 2024	DS DP			
	8691185	Jan 24, 2023	U-336			
	8916131	Sep 16, 2028	DP			

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APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>FLUTICASONE FUROATE - FLONASE SENSIMIST ALLERGY RELIEF</u>						
N 022051	002	6858596	Aug 03, 2021	DP	U-1890	
		7101866	Aug 03, 2021	DS DP	U-1890	
		7541350	Aug 03, 2021	DP	U-1890	
		8062264	Apr 05, 2026	DP		
		8147461	Oct 15, 2028	DP		
		8347879	Jul 15, 2028	DP		
		8752543	Apr 05, 2026	DP		
		9320862	Nov 06, 2024	DP		
<u>FLUTICASONE FUROATE - ARNUITY ELLIPTA</u>						
N 205625	001	7101866	Aug 03, 2021	DS DP	U-1559	NPP May 17, 2021
		7629335	Aug 03, 2021	DP		
		8113199	Oct 23, 2027	DP		
		8161968	Feb 05, 2028	DP		
		8201556	Feb 05, 2029	DP		
		8534281	Mar 08, 2030	DP		
		8746242	Oct 11, 2030	DP		
		9333310	Oct 02, 2027	DP		
<u>FLUTICASONE FUROATE - ARNUITY ELLIPTA</u>						
N 205625	002	7101866	Aug 03, 2021	DS DP	U-1559	
		7629335	Aug 03, 2021	DP		
		8113199	Oct 23, 2027	DP		
		8161968	Feb 05, 2028	DP		
		8201556	Feb 05, 2029	DP		
		8534281	Mar 08, 2030	DP		
		8746242	Oct 11, 2030	DP		
		9333310	Oct 02, 2027	DP		
<u>FLUTICASONE FUROATE - ARNUITY ELLIPTA</u>						
N 205625	003	7101866	Aug 03, 2021	DS DP	U-2349	NS May 17, 2021
		7629335	Aug 03, 2021	DP		
		8113199	Oct 23, 2027	DP		
		8161968	Feb 05, 2028	DP		
		8201556	Feb 05, 2029	DP		
		8534281	Mar 08, 2030	DP		
		8746242	Oct 11, 2030	DP		
		9333310	Oct 02, 2027	DP		
<u>FLUTICASONE FUROATE; UMECLIDINIUM BROMIDE; VILANTEROL TRIFENATATE - TRELEGY ELLIPTA</u>						
N 209482	001	6537983	Aug 03, 2021	DP	U-2125	I-775 Apr 24, 2021
		6759398	Aug 03, 2021	DP	U-2125	
		6878698	Aug 03, 2021		U-2134	
		7101866	Aug 03, 2021	DS DP	U-2126	
		7439393	May 21, 2025	DS DP	U-2127	
		7488827	Dec 18, 2027	DS DP		
		7498440	Apr 27, 2025	DS DP		
		7629335	Aug 03, 2021	DP		
		7776895	Sep 11, 2022	DP		
		8113199	Oct 23, 2027	DP		
		8161968	Feb 05, 2028	DP		
		8183257	Jul 27, 2025		U-2128	
		8309572	Apr 27, 2025		U-2129	
		8511304	Jun 14, 2027	DP		
		8534281	Mar 08, 2030	DP		
		8746242	Oct 11, 2030	DP		
		9333310	Oct 02, 2027	DP		
		9750726	Nov 29, 2030	DP		
		RE44874	Mar 23, 2023	DS DP	U-2127	
<u>FLUTICASONE FUROATE; VILANTEROL TRIFENATATE - BREO ELLIPTA</u>						
N 204275	001	6537983	Aug 03, 2021	DP	U-1401	M-202 May 15, 2020
		6537983	Aug 03, 2021	DP	U-1691	
		6759398	Aug 03, 2021	DP	U-1401	
		6759398	Aug 03, 2021	DP	U-1691	
		6878698	Aug 03, 2021		U-1401	
		7101866	Aug 03, 2021	DS DP	U-1401	
		7101866	Aug 03, 2021	DS DP	U-1691	
		7439393	May 21, 2025	DS DP	U-1401	
		7439393	May 21, 2025	DS DP	U-1691	

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<u>FLUTICASONE FUROATE; VILANTEROL TRIFENATATE - BREO ELLIPTA</u>						
N 204275 001	7439393	May 21, 2025	DS DP U-2099			
	7439393	May 21, 2025	DS DP U-2100			
	7629335	Aug 03, 2021	DP			
	7776895	Sep 11, 2022	DP			
	8113199	Oct 23, 2027	DP			
	8161968	Feb 05, 2028	DP			
	8511304	Jun 14, 2027	DP U-1424			
	8511304	Jun 14, 2027	DP U-1691			
	8534281	Mar 08, 2030	DP			
	8746242	Oct 11, 2030	DP			
	9333310	Oct 02, 2027	DP			
	RE44874	Mar 23, 2023	DS DP U-1548			
	RE44874	Mar 23, 2023	DS DP U-1691			
<u>FLUTICASONE FUROATE; VILANTEROL TRIFENATATE - BREO ELLIPTA</u>						
N 204275 002	6537983	Aug 03, 2021	DP U-1691		M-202	May 15, 2020
	6759398	Aug 03, 2021	DP U-1691			
	7101866	Aug 03, 2021	DS DP U-1691			
	7439393	May 21, 2025	DS DP U-1691			
	7439393	May 21, 2025	DS DP U-2099			
	7439393	May 21, 2025	DS DP U-2100			
	7629335	Aug 03, 2021	DP			
	7776895	Sep 11, 2022	DP			
	8113199	Oct 23, 2027	DP			
	8161968	Feb 05, 2028	DP			
	8511304	Jun 14, 2027	DP U-1691			
	8534281	Mar 08, 2030	DP			
	8746242	Oct 11, 2030	DP			
	9333310	Oct 02, 2027	DP			
	RE44874	Mar 23, 2023	DS DP U-1691			
<u>FLUTICASONE PROPIONATE - FLOVENT HFA</u>						
N 021433 001	7500444	Feb 26, 2026	DP			
	7500444*PED	Aug 26, 2026				
	7832351	Jun 19, 2023	DP			
	9861771	Oct 11, 2020	DP			
<u>FLUTICASONE PROPIONATE - FLOVENT HFA</u>						
N 021433 002	6743413	Jun 01, 2021	U-581	Y		
	7500444	Feb 26, 2026	DP			
	7500444*PED	Aug 26, 2026				
	7832351	Jun 19, 2023	DP			
	9861771	Oct 11, 2020	DP			
<u>FLUTICASONE PROPIONATE - FLOVENT HFA</u>						
N 021433 003	7500444	Feb 26, 2026	DP			
	7500444*PED	Aug 26, 2026				
	7832351	Jun 19, 2023	DP			
	9861771	Oct 11, 2020	DP			
<u>FLUTICASONE PROPIONATE - ARMONAIR RESPICLICK</u>						
N 208798 001	10022510	May 18, 2031	DP		NP	Jan 27, 2020
	10124131	May 18, 2031	DP			
	10195375	Feb 14, 2031	DP			
	6701917	Jun 23, 2021	DP			
	6718972	Jun 23, 2021	DP			
	6748947	Jun 23, 2021	DP			
	6871646	Jun 23, 2021	DP			
	7540282	May 06, 2023	DP			
	8006690	Jun 23, 2021	DP			
	8651103	Mar 26, 2028	DP			
	8714149	Feb 25, 2032	DP			
	8978966	Jan 13, 2032	DP			
	9216260	Jun 28, 2031	DP			
	9463288	May 19, 2025	DP			
	9616024	Sep 01, 2024	DP			
	9731087	May 18, 2031	DP			

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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 002	10022510	May 18, 2031	DP		NP	Jan 27, 2020
	10124131	May 18, 2031	DP			
	10195375	Feb 14, 2031	DP			
	6701917	Jun 23, 2021	DP			
	6718972	Jun 23, 2021	DP			
	6748947	Jun 23, 2021	DP			
	6871646	Jun 23, 2021	DP			
	7540282	May 06, 2023	DP			
	8006690	Jun 23, 2021	DP			
	8651103	Mar 26, 2028	DP			
	8714149	Feb 25, 2032	DP			
	8978966	Jan 13, 2032	DP			
	9216260	Jun 28, 2031	DP			
	9463288	May 19, 2025	DP			
	9616024	Sep 01, 2024	DP			
	9731087	May 18, 2031	DP			
<u>FLUTICASONE PROPIONATE - ARMONAIR RESPICLICK</u>						
N 208798 003	10022510	May 18, 2031	DP		NP	Jan 27, 2020
	10124131	May 18, 2031	DP			
	10195375	Feb 14, 2031	DP			
	6701917	Jun 23, 2021	DP			
	6718972	Jun 23, 2021	DP			
	6748947	Jun 23, 2021	DP			
	6871646	Jun 23, 2021	DP			
	7540282	May 06, 2023	DP			
	8006690	Jun 23, 2021	DP			
	8651103	Mar 26, 2028	DP			
	8714149	Feb 25, 2032	DP			
	8978966	Jan 13, 2032	DP			
	9216260	Jun 28, 2031	DP			
	9463288	May 19, 2025	DP			
	9616024	Sep 01, 2024	DP			
	9731087	May 18, 2031	DP			
<u>FLUTICASONE PROPIONATE - XHANCE</u>						
N 209022 001	10076614	Oct 20, 2034	DP		NP	Sep 18, 2020
	10076615	Jul 30, 2029	U-2133			
	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			
	6715485	Mar 03, 2020	DP			
	7975690	Dec 29, 2025	U-2133			
	8327844	Oct 08, 2023	U-2133			
	8522778	May 11, 2022	DP			
	8550073	Oct 22, 2029	DP			
	8555878	Mar 20, 2020	DP			
	8978647	Aug 06, 2030	DP			
	9072857	Apr 10, 2021	DP			
	9468727	Jul 30, 2020	DP			
<u>FLUTICASONE PROPIONATE; SALMETEROL XINAFOATE - ADVAIR DISKUS 100/50</u>						
N 021077 001					M-214	Dec 20, 2020
<u>FLUTICASONE PROPIONATE; SALMETEROL XINAFOATE - ADVAIR DISKUS 250/50</u>						
N 021077 002					M-214	Dec 20, 2020
<u>FLUTICASONE PROPIONATE; SALMETEROL XINAFOATE - ADVAIR DISKUS 500/50</u>						
N 021077 003					M-214	Dec 20, 2020
<u>FLUTICASONE PROPIONATE; SALMETEROL XINAFOATE - ADVAIR HFA</u>						
N 021254 001	7500444	Feb 26, 2026	DP			
	7500444*PED	Aug 26, 2026				
	7832351	Jun 19, 2023	DP			
	9861771	Oct 11, 2020	DP			

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APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>FLUTICASONE PROPIONATE; SALMETEROL XINAFOATE - ADVAIR HFA</u>						
N 021254	002	7500444	Feb 26, 2026	DP		
		7500444*PED	Aug 26, 2026			
		7832351	Jun 19, 2023	DP		
		9861771	Oct 11, 2020	DP		
<u>FLUTICASONE PROPIONATE; SALMETEROL XINAFOATE - ADVAIR HFA</u>						
N 021254	003	7500444	Feb 26, 2026	DP		
		7500444*PED	Aug 26, 2026			
		7832351	Jun 19, 2023	DP		
		9861771	Oct 11, 2020	DP		
<u>FLUTICASONE PROPIONATE; SALMETEROL XINAFOATE - AIRDUO RESPICLICK</u>						
N 208799	001	10022510	May 18, 2031	DP	NP	Jan 27, 2020
		10124131	May 18, 2031	DP		
		10195375	Feb 14, 2031	DP		
		6701917	Jun 23, 2021	DP		
		6718972	Jun 23, 2021	DP		
		6748947	Jun 23, 2021	DP		
		6871646	Jun 23, 2021	DP		
		7540282	May 06, 2023	DP		
		8006690	Jun 23, 2021	DP		
		8651103	Mar 26, 2028	DP		
		8714149	Feb 25, 2032	DP		
		8978966	Jan 13, 2032	DP		
		9066957	Oct 06, 2034	DP U-645		
		9216260	Jun 28, 2031	DP		
		9415008	Oct 06, 2034	DP U-645		
		9463288	May 19, 2025	DP		
		9616024	Sep 01, 2024	DP		
		9731087	May 18, 2031	DP		
		9987229	Sep 01, 2024	DP		
<u>FLUTICASONE PROPIONATE; SALMETEROL XINAFOATE - AIRDUO RESPICLICK</u>						
N 208799	002	10022510	May 18, 2031	DP	NP	Jan 27, 2020
		10124131	May 18, 2031	DP		
		10195375	Feb 14, 2031	DP		
		6701917	Jun 23, 2021	DP		
		6718972	Jun 23, 2021	DP		
		6748947	Jun 23, 2021	DP		
		6871646	Jun 23, 2021	DP		
		7540282	May 06, 2023	DP		
		8006690	Jun 23, 2021	DP		
		8651103	Mar 26, 2028	DP		
		8714149	Feb 25, 2032	DP		
		8978966	Jan 13, 2032	DP		
		9066957	Oct 06, 2034	DP U-645		
		9216260	Jun 28, 2031	DP		
		9463288	May 19, 2025	DP		
		9616024	Sep 01, 2024	DP		
		9731087	May 18, 2031	DP		
		9987229	Sep 01, 2024	DP		
<u>FLUTICASONE PROPIONATE; SALMETEROL XINAFOATE - AIRDUO RESPICLICK</u>						
N 208799	003	10022510	May 18, 2031	DP	NP	Jan 27, 2020
		10124131	May 18, 2031	DP		
		10195375	Feb 14, 2031	DP		
		6701917	Jun 23, 2021	DP		
		6718972	Jun 23, 2021	DP		
		6748947	Jun 23, 2021	DP		
		6871646	Jun 23, 2021	DP		
		7540282	May 06, 2023	DP		
		8006690	Jun 23, 2021	DP		
		8651103	Mar 26, 2028	DP		
		8714149	Feb 25, 2032	DP		
		8978966	Jan 13, 2032	DP		
		9066957	Oct 06, 2034	DP U-645		
		9216260	Jun 28, 2031	DP		
		9463288	May 19, 2025	DP		
		9616024	Sep 01, 2024	DP		
		9731087	May 18, 2031	DP		

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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	9987229	Sep 01, 2024	DP		
<u>FLUTICASONE PROPIONATE; SALMETEROL XINAFOATE - AIRDUO DIGIHALER</u>						
N 208799	004	10022510	May 18, 2031	DP		
		10124131	May 18, 2031	DP		
		10195375	Feb 14, 2031	DP		
		6701917	Jun 23, 2021	DP		
		6718972	Jun 23, 2021	DP		
		6748947	Jun 23, 2021	DP		
		6871646	Jun 23, 2021	DP		
		7540282	May 06, 2023	DP		
		8006690	Jun 23, 2021	DP		
		8651103	Mar 26, 2028	DP		
		8714149	Feb 25, 2032	DP		
		8978966	Jan 13, 2032	DP		
		9066957	Oct 06, 2034	DP	U-645	
		9216260	Jun 28, 2031	DP		
		9415008	Oct 06, 2034	DP	U-645	
		9463288	May 19, 2025	DP		
		9616024	Sep 01, 2024	DP		
		9731087	May 18, 2031	DP		
		9782550	Aug 28, 2035	DP		
		9782551	Aug 28, 2035	DP		
		9987229	Sep 01, 2024	DP		
<u>FLUTICASONE PROPIONATE; SALMETEROL XINAFOATE - AIRDUO DIGIHALER</u>						
N 208799	005	10022510	May 18, 2031	DP		
		10124131	May 18, 2031	DP		
		10195375	Feb 14, 2031	DP		
		6701917	Jun 23, 2021	DP		
		6718972	Jun 23, 2021	DP		
		6748947	Jun 23, 2021	DP		
		6871646	Jun 23, 2021	DP		
		7540282	May 06, 2023	DP		
		8006690	Jun 23, 2021	DP		
		8651103	Mar 26, 2028	DP		
		8714149	Feb 25, 2032	DP		
		8978966	Jan 13, 2032	DP		
		9066957	Oct 06, 2034	DP	U-645	
		9216260	Jun 28, 2031	DP		
		9463288	May 19, 2025	DP		
		9616024	Sep 01, 2024	DP		
		9731087	May 18, 2031	DP		
		9782550	Aug 28, 2035	DP		
		9782551	Aug 28, 2035	DP		
		9987229	Sep 01, 2024	DP		
<u>FLUTICASONE PROPIONATE; SALMETEROL XINAFOATE - AIRDUO DIGIHALER</u>						
N 208799	006	10022510	May 18, 2031	DP		
		10124131	May 18, 2031	DP		
		10195375	Feb 14, 2031	DP		
		6701917	Jun 23, 2021	DP		
		6718972	Jun 23, 2021	DP		
		6748947	Jun 23, 2021	DP		
		6871646	Jun 23, 2021	DP		
		7540282	May 06, 2023	DP		
		8006690	Jun 23, 2021	DP		
		8651103	Mar 26, 2028	DP		
		8714149	Feb 25, 2032	DP		
		8978966	Jan 13, 2032	DP		
		9066957	Oct 06, 2034	DP	U-645	
		9216260	Jun 28, 2031	DP		
		9463288	May 19, 2025	DP		
		9616024	Sep 01, 2024	DP		
		9731087	May 18, 2031	DP		
		9782550	Aug 28, 2035	DP		
		9782551	Aug 28, 2035	DP		
		9987229	Sep 01, 2024	DP		

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<u>FLUVASTATIN SODIUM - LESCOL XL</u>						
N 021192	001	6242003				
<u>FLUVOXAMINE MALEATE - LUVOX CR</u>						
N 022033	001	7465462				
<u>FLUVOXAMINE MALEATE - LUVOX CR</u>						
N 022033	002	7465462				
<u>FOLLITROPIN ALFA/BETA - GONAL-F RFF REDI-JECT</u>						
N 021684	001	7741268				
<u>FOLLITROPIN ALFA/BETA - GONAL-F RFF REDI-JECT</u>						
N 021684	002	7741268				
<u>FOLLITROPIN ALFA/BETA - GONAL-F RFF REDI-JECT</u>						
N 021684	003	7741268				
<u>FOMEPIZOLE - ANTIZOL</u>						
N 020696	001	7553863				
<u>FORMOTEROL FUMARATE - FORADIL</u>						
N 020831	001	6887459				
<u>FORMOTEROL FUMARATE - PERFOROMIST</u>						
N 022007	001	6667344				
		6814953				
		7348362				
		7462645				
		8623922				
		9730890				
<u>FORMOTEROL FUMARATE; GLYCOPYRROLATE - BEVESPI AEROSPHERE</u>						
N 208294	001	8324266				
		8703806				
		8808713				
		8815258				
		9415009				
		9463161				
<u>FORMOTEROL FUMARATE; MOMETASONE FUROATE - DULERA</u>						
N 022518	001	7067502				
		7067502*PED				
		7566705				
		7566705*PED				
<u>FORMOTEROL FUMARATE; MOMETASONE FUROATE - DULERA</u>						
N 022518	002	7067502				
		7067502*PED				
		7566705				
		7566705*PED				
<u>FOSAPREPITANT DIMEGLUMINE - EMEND</u>						
N 022023	001					
<u>FOSAPREPITANT DIMEGLUMINE - EMEND</u>						
N 022023	002					
<u>FOSNETUPITANT CHLORIDE HYDROCHLORIDE; PALONOSETRON HYDROCHLORIDE - AKYNZEO</u>						
N 210493	001	10208073				
		8426450				
		8895586				
		9186357				
		9403772				
		9908907				
<u>FOSPROPOFOL DISODIUM - LUSEDRA</u>						
N 022244	001	6204257				

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<u>FOSTAMATINIB DISODIUM - TAVALISSE</u>						
N 209299 001	7449458	Sep 04, 2026	DS		NCE	Apr 17, 2023
	7538108	Mar 28, 2026	DS	U-2294	ODE-174	Apr 17, 2025
	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		
	9737554	Jan 19, 2026	DP			
<u>FOSTAMATINIB DISODIUM - TAVALISSE</u>						
N 209299 002	7449458	Sep 04, 2026	DS		NCE	Apr 17, 2023
	7538108	Mar 28, 2026	DS	U-2294	ODE-174	Apr 17, 2025
	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		
	9737554	Jan 19, 2026	DP			
<u>FULVESTRANT - FASLODEX</u>						
N 021344 001	6774122	Jan 09, 2021		U-1826	I-749	Aug 25, 2020
	6774122	Jan 09, 2021		U-2108		
	6774122	Jan 09, 2021		U-2163		
	6774122	Jan 09, 2021		U-2504		
	6774122	Jan 09, 2021		U-596		
	6774122*PED	Jul 09, 2021				
	7456160	Jan 09, 2021		U-1826		
	7456160	Jan 09, 2021		U-2108		
	7456160	Jan 09, 2021		U-2163		
	7456160	Jan 09, 2021		U-2504		
	7456160	Jan 09, 2021		U-596		
	7456160*PED	Jul 09, 2021				
	8329680	Jan 09, 2021		U-1826		
	8329680	Jan 09, 2021		U-2108		
	8329680	Jan 09, 2021		U-2163		
	8329680	Jan 09, 2021		U-2504		
	8329680	Jan 09, 2021		U-596		
	8329680*PED	Jul 09, 2021				
	8466139	Jan 09, 2021		U-1826		
	8466139	Jan 09, 2021		U-2108		
	8466139	Jan 09, 2021		U-2163		
	8466139	Jan 09, 2021		U-2504		
	8466139	Jan 09, 2021		U-596		
	8466139*PED	Jul 09, 2021				
<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>GABAPENTIN - NEURONTIN</u>						
N 021129 001	7256216	May 28, 2022	DP			
<u>GABAPENTIN - GRALISE</u>						
N 022544 001	6488962	Jun 20, 2020	DP			
	6723340	Oct 25, 2021	DP			
	7438927	Feb 26, 2024		U-1114		
	7731989	Oct 25, 2022	DP			

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<u>GABAPENTIN - GRALISE</u>						
N 022544	001	8192756	Oct 25, 2022	DP U-1114		
		8252332	Oct 25, 2022	DP U-1114		
		8333992	Oct 25, 2022	DP U-1114		
<u>GABAPENTIN - GRALISE</u>						
N 022544	002	6488962	Jun 20, 2020	DP		
		6723340	Oct 25, 2021	DP		
		7438927	Feb 26, 2024	U-1114		
		7731989	Oct 25, 2022	DP		
		8192756	Oct 25, 2022	DP U-1114		
		8252332	Oct 25, 2022	DP U-1114		
		8333992	Oct 25, 2022	DP U-1114		
<u>GABAPENTIN ENACARBIL - HORIZANT</u>						
N 022399	001	6818787	Apr 06, 2025	DS DP		
		8026279	Nov 10, 2026	DS DP		
		8048917	Nov 06, 2022	DS DP U-1247		
		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		
		8048917	Nov 06, 2022	DS DP U-1247		
		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>GADOBUTROL - GDAVIST</u>						
N 201277	001				I-801	Jul 12, 2022
<u>GADOBUTROL - GDAVIST</u>						
N 201277	002				I-801	Jul 12, 2022
<u>GADOBUTROL - GDAVIST</u>						
N 201277	003				I-801	Jul 12, 2022
<u>GADOBUTROL - GDAVIST</u>						
N 201277	004				I-801	Jul 12, 2022
<u>GADOBUTROL - GDAVIST</u>						
N 201277	005				I-801	Jul 12, 2022
<u>GADOBUTROL - GDAVIST</u>						
N 201277	006	5980864	Nov 09, 2021	DS DP U-1119	I-801	Jul 12, 2022
<u>GADOFOSVESET TRISODIUM - ABLAVAR</u>						
N 021711	001	6676929	May 04, 2020	DP		
<u>GADOFOSVESET TRISODIUM - ABLAVAR</u>						
N 021711	002	6676929	May 04, 2020	DP		
<u>GADOTERATE MEGLUMINE - DOTAREM</u>						
N 204781	001				NPP	Aug 25, 2020
<u>GADOTERATE MEGLUMINE - DOTAREM</u>						
N 204781	002				NPP	Aug 25, 2020
<u>GADOTERATE MEGLUMINE - DOTAREM</u>						
N 204781	003				NPP	Aug 25, 2020
<u>GADOTERATE MEGLUMINE - DOTAREM</u>						
N 204781	004				NPP	Aug 25, 2020

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<u>GADOTERATE MEGLUMINE - DOTAREM</u>						
N 204781	005				NPP	Aug 25, 2020
<u>GADOXETATE DISODIUM - EOVI</u>						
N 022090	001	6039931	Nov 13, 2021	U-1239		
<u>GADOXETATE DISODIUM - EOVI</u>						
N 022090	002	6039931	Nov 13, 2021	U-1239		
<u>GALLIUM DOTATATE GA-68 - NETSPOT</u>						
N 208547	001	9375498	Aug 10, 2032	DP	NCE ODE-120	Jun 01, 2021 Jun 01, 2023
<u>GALLIUM DOTATOC GA-68 - GALLIUM DOTATOC GA 68</u>						
N 210828	001				NCE W	Aug 21, 2024 Aug 21, 2024
<u>GANCICLOVIR - GANZYK-RTU</u>						
N 209347	001	9486530	Sep 02, 2034	DP		
<u>GATIFLOXACIN - ZYMAR</u>						
N 021493	001	6333045*PED	Feb 20, 2020			
<u>GEFITINIB - IRESSA</u>						
N 206995	001				ODE-95	Jul 13, 2022
<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>GILTERITINIB FUMARATE - XOSPATA</u>						
N 211349	001	8969336	Jan 27, 2031	DS DP	NCE	Nov 28, 2023
		9487491	Jul 28, 2030		ODE-222	Nov 28, 2025
				U-2456		
<u>GIVOSIRAN SODIUM - GIVLAARI</u>						
N 212194	001	10119143	Oct 03, 2034	DS DP	NCE	Nov 20, 2024
		10125364	Mar 15, 2033	DS DP	ODE-273	Nov 20, 2026
		10131907	Aug 24, 2028	DS DP		
		10273477	Mar 08, 2024	DS		
		8106022	Dec 12, 2029	DS DP		
		8546143	Jan 09, 2022	DS		
		8828956	Dec 04, 2028	DS DP		
		9133461	May 14, 2033	DS DP		
		9150605	Aug 28, 2025	DS DP		
		9631193	Mar 15, 2033			
		9708610	Jan 01, 2024	DS DP		

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<u>GIVOSIRAN SODIUM - GIVLAARI</u>						
N 212194	001 9708615	Mar 08, 2024	DS			
<u>GLASDEGIB MALEATE - DAURISMO</u>						
N 210656	001 10414748	Apr 13, 2036	DS DP		NCE	Nov 21, 2023
	8148401	Jan 30, 2031	DS DP		ODE-224	Nov 21, 2025
	8431597	Jun 29, 2028	DP			
<u>GLASDEGIB MALEATE - DAURISMO</u>						
N 210656	002 10414748	Apr 13, 2036	DS DP		NCE	Nov 21, 2023
	8148401	Jan 30, 2031	DS DP		ODE-224	Nov 21, 2025
	8431597	Jun 29, 2028	DP			
<u>GLECAPREVIR; PIBRENTASVIR - MAVYRET</u>						
N 209394	001 10028937	Jun 10, 2030		U-2141	D-175	Sep 26, 2022
	10028937	Jun 10, 2030		U-2532	M-230	Aug 06, 2021
	10039754	Jun 10, 2030		U-2141	NCE	Aug 03, 2022
	10039754	Jun 10, 2030		U-2532	NPP	Apr 30, 2022
	10286029	Mar 14, 2034		U-2532	ODE-232	Apr 30, 2026
	8648037	Jan 19, 2032	DS DP	U-2141	ODE-233	Apr 30, 2026
	8648037	Jan 19, 2032	DS DP	U-2532		
	8937150	May 18, 2032	DS DP			
	9321807	Jun 05, 2035	DS			
	9586978	Jun 10, 2030		U-2141		
	9586978	Jun 10, 2030		U-2532		
<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>GLIMEPIRIDE; ROSIGLITAZONE MALEATE - AVANDARYL</u>						
N 021700	001 7358366	Apr 19, 2020	DS		Y	
	7358366*PED	Oct 19, 2020				
<u>GLIMEPIRIDE; ROSIGLITAZONE MALEATE - AVANDARYL</u>						
N 021700	002 7358366	Apr 19, 2020	DS		Y	
	7358366*PED	Oct 19, 2020				
<u>GLIMEPIRIDE; ROSIGLITAZONE MALEATE - AVANDARYL</u>						
N 021700	003 7358366	Apr 19, 2020	DS		Y	
	7358366*PED	Oct 19, 2020				
<u>GLIMEPIRIDE; ROSIGLITAZONE MALEATE - AVANDARYL</u>						
N 021700	004 7358366	Apr 19, 2020	DS		Y	
	7358366*PED	Oct 19, 2020				
<u>GLIMEPIRIDE; ROSIGLITAZONE MALEATE - AVANDARYL</u>						
N 021700	005 7358366	Apr 19, 2020	DS		Y	
	7358366*PED	Oct 19, 2020				
<u>GLUCAGON - BAOSIMI</u>						
N 210134	001 10213487	Feb 16, 2036	DP	U-2604	NP	Jul 24, 2022
	6938798	Jan 03, 2022	DP			
<u>GLUCAGON - GVOKE PFS</u>						
N 212097	001 9649364	Apr 22, 2036	DP	U-2742	NP	Sep 10, 2022
<u>GLUCAGON - GVOKE PFS</u>						
N 212097	002 9649364	Apr 22, 2036	DP	U-2742	NP	Sep 10, 2022
<u>GLUCAGON - GVOKE HYPOPEN</u>						
N 212097	003 9649364	Apr 22, 2036	DP	U-2742	NP	Sep 10, 2022
<u>GLUCAGON - GVOKE HYPOPEN</u>						
N 212097	004 9649364	Apr 22, 2036	DP	U-2742	NP	Sep 10, 2022

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<u>GLYCEROL PHENYLBUTYRATE - RAVICTI</u>						
N 203284 001	10045958	Sep 22, 2030	U-1816		NPP	Apr 28, 2020
	10045959	Sep 22, 2030	U-1816		ODE-157	Apr 28, 2024
	10183002	Sep 22, 2030	U-1816		ODE-42	Feb 01, 2020
	10183003	Sep 22, 2030	U-1816			
	10183004	Sep 22, 2030	U-1816			
	10183005	Sep 22, 2030	U-1816			
	10183006	Sep 22, 2030	U-1816			
	8404215	Mar 09, 2032	U-1383			
	8642012	Sep 22, 2030	U-1383			
	9095559	Mar 09, 2032	U-1383			
	9254278	Mar 09, 2032	U-1816			
	9326966	Mar 09, 2032	U-1816			
	9561197	Sep 22, 2030	U-1383			
	9962359	Sep 22, 2030	U-1816			
	9999608	Sep 22, 2030	U-1816			
<u>GLYCOPYRROLATE - CUVPOSA</u>						
N 022571 001	7638552	Aug 20, 2023	U-1076			
	7816396	Aug 20, 2023	U-1076			
<u>GLYCOPYRROLATE - SEEBRI</u>						
N 207923 001	7229607	Apr 09, 2021	U-1773			
	7736670	Jun 27, 2021	DP			
	8029768	Apr 09, 2021	U-1773			
	8048451	Jun 27, 2021	DP			
	8182838	Oct 20, 2028	DP			
	8303991	Jun 27, 2021	DP			
	8435567	Jun 27, 2021	DP			
	8479730	Oct 11, 2028	DP			
	8580306	Jun 27, 2021	DP			
	8956661	Jun 27, 2021	DP			
	9931304	Jun 27, 2021	DP			
	9962338	Jun 27, 2021	DP			
<u>GLYCOPYRROLATE - LONHALA MAGNAIR KIT</u>						
N 208437 001	10376661	Sep 14, 2035	DP		NP	Dec 05, 2020
	6962151	Oct 27, 2020	DP			
	7316067	Sep 06, 2022	DP			
	7458372	Nov 18, 2024	DP			
	7931212	Nov 25, 2025	DP			
	8511581	Nov 08, 2023	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			
	7229607	Apr 09, 2021	U-1773			
	7736670	Jun 27, 2021	DP			
	7820694	Jun 02, 2020	DP U-1773			
	8029768	Apr 09, 2021	U-1773			
	8048451	Jun 27, 2021	DP			
	8067437	Jun 02, 2020	U-1773			
	8182838	Oct 20, 2028	DP			
	8283362	Jun 02, 2020	DP U-1773			
	8303991	Jun 27, 2021	DP			
	8435567	Jun 27, 2021	DP			
	8479730	Oct 11, 2028	DP			
	8580306	Jun 27, 2021	DP			
	8658673	Jun 02, 2020	DP U-1773			
	8796307	Jun 02, 2020	DP			
	8956661	Jun 27, 2021	DP			
	9931304	Jun 27, 2021	DP			
	9962338	Jun 27, 2021	DP			
<u>GLYCOPYRROLATE; TOSYLATE - OBREXZA</u>						
N 210361 001	10004717	Feb 28, 2033	DP U-2398			
	10052267	Oct 17, 2028	DP U-2398			
	10543192	Feb 28, 2033	DP			

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<u>GLYCOPYRRONIUM TOSYLATE - QBREXZA</u>						
N 210361	001	10548875	Feb 28, 2033	DS DP	U-2398	
		6433003	Apr 10, 2020		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		NCE Dec 12, 2024
		10266827	Jun 28, 2025		U-2675	ODE-280 Dec 12, 2026
		10421966	Jun 28, 2025	DS DP		
		10533174	May 04, 2021	DP		
		9024007	Jun 28, 2025	DS DP		
		9416361	May 04, 2021	DS		
		9994851	Jun 28, 2025	DS DP		
		RE47691	Jun 28, 2025	DP		
<u>GOSERELIN ACETATE - ZOLADEX</u>						
N 019726	001	7118552	Apr 13, 2022	DP		
		7220247	Apr 09, 2022	DP		
		7500964	Feb 26, 2021	DP		
<u>GOSERELIN ACETATE - ZOLADEX</u>						
N 020578	001	7118552	Apr 13, 2022	DP		
		7220247	Apr 09, 2022	DP		
		7500964	Feb 26, 2021	DP		
<u>GRANISETRON - SANCUSO</u>						
N 022198	001	7608282	Jan 22, 2025	DP	U-1011	
<u>GRANISETRON - SUSTOL</u>						
N 022445	001	10357570	Sep 28, 2024		U-2253	
		6613355	Jun 28, 2021	DP		
		6790458	May 11, 2021	DP		
		8252304	Sep 28, 2024	DP		
		8252305	Sep 28, 2024		U-1891	
		8715710	Sep 28, 2024	DP		
		9913910	Sep 28, 2024		U-2253	
<u>GUAIFENESIN - MUCINEX</u>						
N 021282	001	6372252	Apr 28, 2020		U-489	
		6955821	Apr 28, 2020	DP	U-489	
		7838032	Apr 28, 2020	DP		
<u>GUAIFENESIN - MUCINEX</u>						
N 021282	002	6372252	Apr 28, 2020		U-489	
		6955821	Apr 28, 2020	DP	U-489	
		7838032	Apr 28, 2020	DP		
<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>GUAIFENESIN; PSEUDOEPHEDRINE HYDROCHLORIDE - MUCINEX D</u>						
N 021585	001	6372252	Apr 28, 2020	DP		
		6955821	Apr 28, 2020	DP	U-686	
		7838032	Apr 28, 2020	DP		
<u>GUAIFENESIN; PSEUDOEPHEDRINE HYDROCHLORIDE - MUCINEX D</u>						
N 021585	002	6372252	Apr 28, 2020	DP		
		6955821	Apr 28, 2020	DP	U-686	
		7838032	Apr 28, 2020	DP		
<u>GUANFACINE HYDROCHLORIDE - INTUNIV</u>						
N 022037	001	6287599	Dec 20, 2020	DP		
		6287599*PED	Jun 20, 2021			
		6811794	Jul 04, 2022	DP	U-494	
		6811794*PED	Jan 04, 2023			

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<u>GUANFACINE HYDROCHLORIDE - INTUNIV</u>						
N 022037	002	6287599	Dec 20, 2020	DP		
		6287599*PED	Jun 20, 2021			
		6811794	Jul 04, 2022	DP U-494		
		6811794*PED	Jan 04, 2023			
<u>GUANFACINE HYDROCHLORIDE - INTUNIV</u>						
N 022037	003	6287599	Dec 20, 2020	DP		
		6287599*PED	Jun 20, 2021			
		6811794	Jul 04, 2022	DP U-494		
		6811794*PED	Jan 04, 2023			
<u>GUANFACINE HYDROCHLORIDE - INTUNIV</u>						
N 022037	004	6287599	Dec 20, 2020	DP		
		6287599*PED	Jun 20, 2021			
		6811794	Jul 04, 2022	DP U-494		
		6811794*PED	Jan 04, 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	NP	Nov 06, 2021
		6517847	Aug 03, 2020	DP		
		8809307	Nov 02, 2031	DP		
<u>HALOBETASOL PROPIONATE - LEXETTE</u>						
N 210566	001				NDF	May 24, 2021
<u>HALOBETASOL PROPIONATE; TAZAROTENE - DUOBRII</u>						
N 209354	001	10251895	Jun 06, 2036	DP	NP	Apr 25, 2022
		10426787	Jun 06, 2036	U-2625		
		10478502	Nov 02, 2031	DP U-2625		
		6517847	Aug 03, 2020	DP		
		8809307	Nov 02, 2031	DP		
<u>HEXAMINOLEVULINATE HYDROCHLORIDE - CYSVIEW KIT</u>						
N 022555	001	10556010	Dec 19, 2036	U-2250	M-220	Feb 15, 2021
		7348361	Nov 06, 2020	DP U-1087		
		7348361	Nov 06, 2020	DP U-2250		
<u>HISTRELIN ACETATE - SUPPRELIN LA</u>						
N 022058	001	8062652	Jun 16, 2026	U-1197		
<u>HYALURONIDASE RECOMBINANT HUMAN - HYLENEX RECOMBINANT</u>						
N 021859	001	7767429	Sep 23, 2027	DS DP		
<u>HYDRALAZINE HYDROCHLORIDE; ISOSORBIDE DINITRATE - BIDIL</u>						
N 020727	001	6465463	Sep 08, 2020	U-71		
		6784177	Sep 08, 2020	U-71		
<u>HYDROCHLOROTHIAZIDE; TELMISARTAN - MICARDIS HCT</u>						
N 021162	001	6358986	Jan 10, 2020			
<u>HYDROCHLOROTHIAZIDE; TELMISARTAN - MICARDIS HCT</u>						
N 021162	002	6358986	Jan 10, 2020			
<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		
		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		

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PRESCRIPTION AND OTC DRUG PRODUCT 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	001	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
		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	003	10028946	Jul 25, 2033			U-1810
		10092559	Sep 12, 2034			U-55
		10322120	Jul 25, 2033			DP
		10456393	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
		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
		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

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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 006	10028946	Jul 25, 2033				U-1810
	10092559	Sep 12, 2034				U-55
	10322120	Jul 25, 2033	DP			
	10456393	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	10130591	Nov 20, 2023	DP			U-1819
	10369109	Jun 16, 2023	DP			
	6733783	Oct 30, 2021	DP			U-1556
	8309060	Nov 20, 2023	DP			U-1556
	8361499	Oct 30, 2021	DP			
	8529948	Aug 06, 2022	DP			
	8551520	Oct 30, 2021	DP			
	8647667	Oct 30, 2021	DP			
	8808740	Dec 21, 2031	DP			U-1556
	9023401	Oct 30, 2020	DP			
	9056052	Oct 30, 2020	DP			
	9060940	Oct 30, 2020				U-1556
	9084816	Aug 24, 2027	DP			
	9095614	Aug 24, 2027				U-1556
	9095615	Aug 24, 2027	DP			
	9198863	Oct 30, 2020	DP			
	9205056	Oct 30, 2020	DP			
	9289391	Oct 30, 2020	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
	9517236	Oct 30, 2020	DP			
	9545380	Aug 24, 2027				U-1556
	9572779	Dec 21, 2031	DP			
	9572804	Oct 30, 2020	DP			
	9669023	Oct 30, 2020	DP			
	9669024	Oct 30, 2020	DP			
	9675610	Jun 16, 2023	DP			
	9675611	Oct 30, 2020				U-1556
	9682077	Oct 30, 2020				U-1556
	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 002	10130591	Nov 20, 2023	DP			U-1819
	10369109	Jun 16, 2023	DP			
	6733783	Oct 30, 2021	DP			U-1556
	8309060	Nov 20, 2023	DP			U-1556
	8361499	Oct 30, 2021	DP			
	8529948	Aug 06, 2022	DP			
	8551520	Oct 30, 2021	DP			
	8647667	Oct 30, 2021	DP			
	8808740	Dec 21, 2031	DP			U-1556
	9023401	Oct 30, 2020	DP			
	9056052	Oct 30, 2020	DP			
	9060940	Oct 30, 2020				U-1556

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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 002	9084816	Aug 24, 2027	DP			
	9095614	Aug 24, 2027		U-1556		
	9095615	Aug 24, 2027	DP			
	9198863	Oct 30, 2020	DP			
	9205056	Oct 30, 2020	DP			
	9289391	Oct 30, 2020	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		
	9517236	Oct 30, 2020	DP			
	9545380	Aug 24, 2027		U-1556		
	9572779	Dec 21, 2031	DP			
	9572804	Oct 30, 2020	DP			
	9669023	Oct 30, 2020	DP			
	9669024	Oct 30, 2020	DP			
	9675610	Jun 16, 2023	DP			
	9675611	Oct 30, 2020		U-1556		
	9682077	Oct 30, 2020		U-1556		
	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	10130591	Nov 20, 2023	DP	U-1819		
	10369109	Jun 16, 2023	DP			
	6733783	Oct 30, 2021	DP	U-1556		
	8309060	Nov 20, 2023	DP	U-1556		
	8361499	Oct 30, 2021	DP			
	8529948	Aug 06, 2022	DP			
	8551520	Oct 30, 2021	DP			
	8647667	Oct 30, 2021	DP			
	8808740	Dec 21, 2031	DP	U-1556		
	9023401	Oct 30, 2020	DP			
	9056052	Oct 30, 2020	DP			
	9060940	Oct 30, 2020		U-1556		
	9084816	Aug 24, 2027	DP			
	9095614	Aug 24, 2027		U-1556		
	9095615	Aug 24, 2027	DP			
	9198863	Oct 30, 2020	DP			
	9205056	Oct 30, 2020	DP			
	9289391	Oct 30, 2020	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		
	9517236	Oct 30, 2020	DP			
	9545380	Aug 24, 2027		U-1556		
	9572779	Dec 21, 2031	DP			
	9572804	Oct 30, 2020	DP			
	9669023	Oct 30, 2020	DP			
	9669024	Oct 30, 2020	DP			
	9675610	Jun 16, 2023	DP			
	9675611	Oct 30, 2020		U-1556		
	9682077	Oct 30, 2020		U-1556		
	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			

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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	10130591	Nov 20, 2023	DP U-1819			
	10369109	Jun 16, 2023	DP			
	6733783	Oct 30, 2021	DP U-1556			
	8309060	Nov 20, 2023	DP U-1556			
	8361499	Oct 30, 2021	DP			
	8529948	Aug 06, 2022	DP			
	8551520	Oct 30, 2021	DP			
	8647667	Oct 30, 2021	DP			
	8808740	Dec 21, 2031	DP U-1556			
	9023401	Oct 30, 2020	DP			
	9056052	Oct 30, 2020	DP			
	9060940	Oct 30, 2020	U-1556			
	9084816	Aug 24, 2027	DP			
	9095614	Aug 24, 2027	U-1556			
	9095615	Aug 24, 2027	DP			
	9198863	Oct 30, 2020	DP			
	9205056	Oct 30, 2020	DP			
	9289391	Oct 30, 2020	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			
	9517236	Oct 30, 2020	DP			
	9545380	Aug 24, 2027	U-1556			
	9572779	Dec 21, 2031	DP			
	9572804	Oct 30, 2020	DP			
	9669023	Oct 30, 2020	DP			
	9669024	Oct 30, 2020	DP			
	9675610	Jun 16, 2023	DP			
	9675611	Oct 30, 2020	U-1556			
	9682077	Oct 30, 2020	U-1556			
	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	10130591	Nov 20, 2023	DP U-1819			
	10369109	Jun 16, 2023	DP			
	6733783	Oct 30, 2021	DP U-1556			
	8309060	Nov 20, 2023	DP U-1556			
	8361499	Oct 30, 2021	DP			
	8529948	Aug 06, 2022	DP			
	8551520	Oct 30, 2021	DP			
	8647667	Oct 30, 2021	DP			
	8808740	Dec 21, 2031	DP U-1556			
	9056052	Oct 30, 2020	DP			
	9060940	Oct 30, 2020	U-1556			
	9084816	Aug 24, 2027	DP			
	9095614	Aug 24, 2027	U-1556			
	9095615	Aug 24, 2027	DP			
	9198863	Oct 30, 2020	DP			
	9205056	Oct 30, 2020	DP			
	9289391	Oct 30, 2020	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			
	9517236	Oct 30, 2020	DP			
	9545380	Aug 24, 2027	U-1556			
	9572779	Dec 21, 2031	DP			
	9572804	Oct 30, 2020	DP			
	9669023	Oct 30, 2020	DP			
	9669024	Oct 30, 2020	DP			
	9675610	Jun 16, 2023	DP			

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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 005	9675611	Oct 30, 2020				U-1556
	9682077	Oct 30, 2020				U-1556
	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	10130591	Nov 20, 2023	DP			U-1819
	10369109	Jun 16, 2023	DP			
	6733783	Oct 30, 2021	DP			U-1556
	8309060	Nov 20, 2023	DP			U-1556
	8361499	Oct 30, 2021	DP			
	8529948	Aug 06, 2022	DP			
	8551520	Oct 30, 2021	DP			
	8647667	Oct 30, 2021	DP			
	8808740	Dec 21, 2031	DP			U-1556
	9056052	Oct 30, 2020	DP			
	9060940	Oct 30, 2020				U-1556
	9084816	Aug 24, 2027	DP			
	9095614	Aug 24, 2027				U-1556
	9095615	Aug 24, 2027	DP			
	9198863	Oct 30, 2020	DP			
	9205056	Oct 30, 2020	DP			
	9289391	Oct 30, 2020	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
	9517236	Oct 30, 2020	DP			
	9545380	Aug 24, 2027				U-1556
	9572779	Dec 21, 2031	DP			
	9572804	Oct 30, 2020	DP			
	9669023	Oct 30, 2020	DP			
	9669024	Oct 30, 2020	DP			
	9675610	Jun 16, 2023	DP			
	9675611	Oct 30, 2020				U-1556
	9682077	Oct 30, 2020				U-1556
	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	10130591	Nov 20, 2023	DP			U-1819
	10369109	Jun 16, 2023	DP			
	6733783	Oct 30, 2021	DP			U-1556
	8309060	Nov 20, 2023	DP			U-1556
	8361499	Oct 30, 2021	DP			
	8529948	Aug 06, 2022	DP			
	8551520	Oct 30, 2021	DP			
	8647667	Oct 30, 2021	DP			
	8808740	Dec 21, 2031	DP			U-1556
	9056052	Oct 30, 2020	DP			
	9060940	Oct 30, 2020				U-1556
	9084816	Aug 24, 2027	DP			
	9095614	Aug 24, 2027				U-1556
	9095615	Aug 24, 2027	DP			
	9198863	Oct 30, 2020	DP			
	9205056	Oct 30, 2020	DP			
	9289391	Oct 30, 2020	DP			
	9486412	Aug 24, 2027	DP			
	9486413	Aug 24, 2027	DP			
	9492389	Aug 24, 2027	DP			

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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	007	9492390	Aug 24, 2027	U-1556		
		9492391	Aug 24, 2027	U-1556		
		9517236	Oct 30, 2020	DP		
		9545380	Aug 24, 2027	U-1556		
		9572779	Dec 21, 2031	DP		
		9572804	Oct 30, 2020	DP		
		9669023	Oct 30, 2020	DP		
		9669024	Oct 30, 2020	DP		
		9675610	Jun 16, 2023	DP		
		9675611	Oct 30, 2020	U-1556		
		9682077	Oct 30, 2020	U-1556		
		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		
<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 BUTYRATE - LOCOID</u>						
N 022076	001	7378405	Dec 19, 2026	DP		
		7981877	Jan 23, 2025	DP		
<u>HYDROGEN PEROXIDE - ESKATA</u>						
N 209305	001	10098910	Apr 21, 2035	DP U-2205	NP	Dec 14, 2020
		10493103	Apr 21, 2035	DP		
		7381427	Jun 08, 2022	U-2205		
		9675639	Jul 04, 2035	DP U-2205		
		9980983	Apr 21, 2035	U-2205		
<u>HYDROMORPHONE HYDROCHLORIDE - DILAUDID-HP</u>						
N 019034	001	6589960	Nov 09, 2020	DP		
		9248229	Mar 12, 2034	DP		
		9731082	Apr 23, 2032	DP		
<u>HYDROMORPHONE HYDROCHLORIDE - DILAUDID-HP</u>						
N 019034	002	6589960	Nov 09, 2020	DP		
		9248229	Mar 12, 2034	DP		
		9731082	Apr 23, 2032	DP		
<u>HYDROMORPHONE HYDROCHLORIDE - DILAUDID</u>						
N 019034	003	6589960	Nov 09, 2020	DS DP		
		9248229	Mar 12, 2034	DP		
		9731082	Apr 23, 2032	DP		

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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	004	6589960	Nov 09, 2020	DS DP		
		9248229	Mar 12, 2034	DP		
		9731082	Apr 23, 2032	DP		
<u>HYDROMORPHONE HYDROCHLORIDE - DILAUDID</u>						
N 019034	005	6589960	Nov 09, 2020	DS DP		
		9248229	Mar 12, 2034	DP		
		9731082	Apr 23, 2032	DP		
<u>HYDROMORPHONE HYDROCHLORIDE - DILAUDID</u>						
N 019891	001	6589960	Nov 09, 2020	DS DP		
<u>HYDROMORPHONE HYDROCHLORIDE - DILAUDID</u>						
N 019892	001	6589960	Nov 09, 2020	DS DP		
<u>HYDROMORPHONE HYDROCHLORIDE - DILAUDID</u>						
N 019892	002	6589960	Nov 09, 2020	DS DP		
<u>HYDROMORPHONE HYDROCHLORIDE - DILAUDID</u>						
N 019892	003	6589960	Nov 09, 2020	DS DP		
<u>HYDROMORPHONE HYDROCHLORIDE - PALLADONE</u>						
N 021044	001	6589960	Nov 09, 2020	DP		
<u>HYDROMORPHONE HYDROCHLORIDE - PALLADONE</u>						
N 021044	002	6589960	Nov 09, 2020	DP		
<u>HYDROMORPHONE HYDROCHLORIDE - PALLADONE</u>						
N 021044	003	6589960	Nov 09, 2020	DP		
<u>HYDROMORPHONE HYDROCHLORIDE - PALLADONE</u>						
N 021044	004	6589960	Nov 09, 2020	DP		
<u>HYDROXYPROGESTERONE CAPROATE - MAKENA (AUTOINJECTOR)</u>						
N 021945	004	10471075	May 02, 2036		U-2236	
		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		
		9789257	Feb 11, 2034	DP		
		9844558	May 02, 2036		U-2236	
<u>HYDROXYUREA - SIKLOS</u>						
N 208843	001				NP ODE-177	Dec 21, 2020 Dec 21, 2024
<u>HYDROXYUREA - SIKLOS</u>						
N 208843	002				NP ODE-177	Dec 21, 2020 Dec 21, 2024
<u>IBANDRONATE SODIUM - BONIVA</u>						
N 021455	002	7192938	May 06, 2023		U-798	
		7410957	May 06, 2023		U-887	
		7718634	May 06, 2023		U-642	
<u>IBRUTINIB - IMBRUVICA</u>						
N 205552	001	10004746	Jun 03, 2031		U-1684	
		10004746	Jun 03, 2031		U-1946	
		10004746	Jun 03, 2031		U-2241	
		10004746	Jun 03, 2031		U-2242	
		10016435	Jun 03, 2031		U-1650	
		10106548	Jun 03, 2033	DS DP	ODE-117	May 06, 2023
		10125140	Jun 03, 2033	DS DP	ODE-128	Jan 18, 2024
		10294231	Jun 03, 2033	DP	ODE-152	Aug 02, 2024
		10294232	Jun 03, 2033	DP	ODE-55	Nov 13, 2020
		10463668	Oct 24, 2034		U-2654	
		10478439	Jun 03, 2031		U-1456	
		10478439	Jun 03, 2031		U-1650	
		10478439	Jun 03, 2031		U-1684	
		10478439	Jun 03, 2031		U-1946	
					D-176	Aug 24, 2021
					I-741	Jan 18, 2020
					I-753	Aug 02, 2020
					M-236	Jan 25, 2022
					ODE-109	Mar 04, 2023
					ODE-117	May 06, 2023
					ODE-128	Jan 18, 2024
					ODE-152	Aug 02, 2024
					ODE-55	Nov 13, 2020
					ODE-60	Feb 12, 2021
					ODE-72	Jul 28, 2021
					ODE-86	Jan 29, 2022

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<u>IBRUTINIB - IMBRUVICA</u>						
N 205552 001	10478439	Jun 03, 2031				
	10478439	Jun 03, 2031				
	10478439	Jun 03, 2031				
	10478439	Jun 03, 2031				
	7514444	Dec 28, 2026	DS DP			
	8008309	Dec 28, 2026	DS DP			
	8476284	Dec 28, 2026			U-1456	
	8476284	Dec 28, 2026			U-1650	
	8476284	Dec 28, 2026			U-1946	
	8476284	Dec 28, 2026			U-1947	
	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	
	8563563	Apr 26, 2027			U-1491	
	8563563	Apr 26, 2027			U-1650	
	8563563	Apr 26, 2027			U-1946	
	8563563	Apr 26, 2027			U-2219	
	8697711	Dec 28, 2026	DS DP			
	8703780	Dec 28, 2026			U-1491	
	8735403	Dec 28, 2026	DS DP			
	8754090	Jun 03, 2031			U-1456	
	8754091	Dec 28, 2026	DP			
	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	
	8957079	Dec 28, 2026	DS DP			
	8999999	Jun 03, 2031			U-1683	
	8999999	Jun 03, 2031			U-1684	
	9125889	Jun 03, 2031			U-1745	
	9181257	Dec 28, 2026	DS DP			
	9296753	Oct 30, 2033	DS DP			
	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	
	9713617	Jun 03, 2033	DP			
	9725455	Jun 03, 2033	DS			
	9795604	Oct 24, 2034			U-2150	
	9801881	Jun 03, 2031			U-1491	
	9801883	Jun 03, 2031			U-2159	
	9814721	Jun 03, 2031			U-1947	
<u>IBRUTINIB - IMBRUVICA</u>						
N 205552 002	10004746	Jun 03, 2031			D-176	Aug 24, 2021
	10004746	Jun 03, 2031			M-236	Jan 25, 2022
	10004746	Jun 03, 2031				
	10004746	Jun 03, 2031				
	10016435	Jun 03, 2031			U-1650	
	10106548	Jun 03, 2033	DS DP			
	10125140	Jun 03, 2033	DS DP			
	10294231	Jun 03, 2033	DP			
	10294232	Jun 03, 2033	DP			
	10463668	Oct 24, 2034			U-2654	
	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	
	7514444	Dec 28, 2026	DS DP			
	8008309	Dec 28, 2026	DS DP			
	8476284	Dec 28, 2026			U-1456	

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<u>IBRUTINIB - IMBRUVICA</u>						
N 205552 002	8476284	Dec 28, 2026				
	8476284	Dec 28, 2026				
	8476284	Dec 28, 2026				
	8497277	Dec 28, 2026				
	8497277	Dec 28, 2026				
	8497277	Dec 28, 2026				
	8497277	Dec 28, 2026				
	8497277	Dec 28, 2026				
	8563563	Apr 26, 2027				
	8563563	Apr 26, 2027				
	8563563	Apr 26, 2027				
	8563563	Apr 26, 2027				
	8697711	Dec 28, 2026	DS DP			
	8703780	Dec 28, 2026				
	8735403	Dec 28, 2026	DS DP			
	8754090	Jun 03, 2031				
	8754091	Dec 28, 2026				
	8952015	Dec 28, 2026				
	8952015	Dec 28, 2026				
	8952015	Dec 28, 2026				
	8952015	Dec 28, 2026				
	8952015	Dec 28, 2026				
	8952015	Dec 28, 2026				
	8957079	Dec 28, 2026	DS DP			
	8999999	Jun 03, 2031				
	8999999	Jun 03, 2031				
	8999999	Jun 03, 2031				
	9125889	Jun 03, 2031				
	9181257	Dec 28, 2026	DS			
	9296753	Oct 30, 2033	DS			
	9540382	Aug 18, 2033				
	9540382	Aug 18, 2033				
	9540382	Aug 18, 2033				
	9540382	Aug 18, 2033				
	9540382	Aug 18, 2033				
	9540382	Aug 18, 2033				
	9713617	Jun 03, 2033				
	9725455	Jun 03, 2033	DS			
	9795604	Oct 24, 2034				
	9801881	Jun 03, 2031				
	9801883	Jun 03, 2031				
	9814721	Jun 03, 2031				
<u>IBRUTINIB - IMBRUVICA</u>						
N 210563 001	10004746	Jun 03, 2031			D-176	Aug 24, 2021
	10004746	Jun 03, 2031			M-236	Jan 25, 2022
	10004746	Jun 03, 2031				
	10004746	Jun 03, 2031				
	10010507	Mar 03, 2036	DP			
	10016435	Jun 03, 2031				
	10106548	Jun 03, 2033	DS DP			
	10125140	Jun 03, 2033	DS DP			
	10213386	Mar 03, 2036	DP			
	10463668	Oct 24, 2034				
	10478439	Jun 03, 2031				
	10478439	Jun 03, 2031				
	10478439	Jun 03, 2031				
	10478439	Jun 03, 2031				
	10478439	Jun 03, 2031				
	10478439	Jun 03, 2031				
	10478439	Jun 03, 2031				
	10478439	Jun 03, 2031				
	10478439	Jun 03, 2031				
	10478439	Jun 03, 2031				
	7514444	Dec 28, 2026	DS DP			
	8008309	Dec 28, 2026	DS DP			
	8476284	Dec 28, 2026				
	8476284	Dec 28, 2026				
	8476284	Dec 28, 2026				
	8476284	Dec 28, 2026				
	8476284	Dec 28, 2026				
	8497277	Dec 28, 2026				
	8497277	Dec 28, 2026				

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<u>IBRUTINIB - IMBRUVICA</u>						
N 210563 001	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			
	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			
	8697711	Dec 28, 2026	DS DP			
	8703780	Dec 28, 2026	U-1491			
	8703780	Dec 28, 2026	U-2242			
	8735403	Dec 28, 2026	DS DP			
	8754090	Jun 03, 2031	U-1456			
	8754091	Dec 28, 2026	DP			
	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			
	8957079	Dec 28, 2026	DS DP			
	8999999	Jun 03, 2031	U-1491			
	8999999	Jun 03, 2031	U-1946			
	8999999	Jun 03, 2031	U-2241			
	8999999	Jun 03, 2031	U-2242			
	9125889	Jun 03, 2031	U-1650			
	9181257	Dec 28, 2026	DS			
	9296753	Oct 30, 2033	DS			
	9655857	Mar 03, 2036	DP			
	9725455	Jun 03, 2033	DS			
	9795604	Oct 24, 2034	U-2150			
	9801881	Jun 03, 2031	U-1491			
	9801881	Jun 03, 2031	U-2242			
	9801883	Jun 03, 2031	U-2159			
	9801883	Jun 03, 2031	U-2243			
	9814721	Jun 03, 2031	U-1947			
<u>IBRUTINIB - IMBRUVICA</u>						
N 210563 002	10004746	Jun 03, 2031	U-1684		D-176	Aug 24, 2021
	10004746	Jun 03, 2031	U-1946		M-236	Jan 25, 2022
	10004746	Jun 03, 2031	U-2241			
	10004746	Jun 03, 2031	U-2242			
	10010507	Mar 03, 2036	DP			
	10016435	Jun 03, 2031	U-1650			
	10106548	Jun 03, 2033	DS DP			
	10125140	Jun 03, 2033	DS DP			
	10213386	Mar 03, 2036	DP			
	10463668	Oct 24, 2034	U-2654			
	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			
	7514444	Dec 28, 2026	DS DP			
	8008309	Dec 28, 2026	DS DP			
	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			
	8497277	Dec 28, 2026	U-1456			
	8497277	Dec 28, 2026	U-1491			
	8497277	Dec 28, 2026	U-1650			

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<u>IBRUTINIB - IMBRUVICA</u>						
N 210563 004	8497277	Dec 28, 2026	U-2241			
	8497277	Dec 28, 2026	U-2242			
	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			
	8697711	Dec 28, 2026	DS DP			
	8703780	Dec 28, 2026	U-1491			
	8703780	Dec 28, 2026	U-2242			
	8735403	Dec 28, 2026	DS DP			
	8754090	Jun 03, 2031	U-1456			
	8754091	Dec 28, 2026	DP			
	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			
	8957079	Dec 28, 2026	DS DP			
	8999999	Jun 03, 2031	U-1491			
	8999999	Jun 03, 2031	U-1946			
	8999999	Jun 03, 2031	U-2241			
	8999999	Jun 03, 2031	U-2242			
	9125889	Jun 03, 2031	U-1650			
	9181257	Dec 28, 2026	DS			
	9296753	Oct 30, 2033	DS			
	9655857	Mar 03, 2036	DP			
	9725455	Jun 03, 2033	DS			
	9795604	Oct 24, 2034	U-2150			
	9801881	Jun 03, 2031	U-1491			
	9801881	Jun 03, 2031	U-2242			
	9801883	Jun 03, 2031	U-2159			
	9801883	Jun 03, 2031	U-2243			
	9814721	Jun 03, 2031	U-1947			
<u>IBUPROFEN - CHILDREN'S ADVIL-FLAVORED</u>						
N 020589 002	10238640	May 25, 2024	DP			
<u>IBUPROFEN - CALDOLOR</u>						
N 022348 001	6727286	Nov 27, 2021	DP U-981			
<u>IBUPROFEN - CALDOLOR</u>						
N 022348 002	6727286	Nov 27, 2021	DP U-981			
	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	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	6342530	Nov 14, 2020	DP U-1127			
	6342530	Nov 14, 2020	DP U-794			
	6344479	Mar 20, 2021	DS DP U-794	Y		
	8415337	Mar 02, 2032	DS DP			
<u>ICOSAPENT ETHYL - VASCEPA</u>						
N 202057 001	10010517	Apr 29, 2030	U-2690		I-819	Dec 13, 2022
	10265287	Apr 29, 2030	U-2700			
	10278935	Jun 28, 2033	U-2701			

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<u>ICOSAPENT ETHYL - VASCEPA</u>						
N 202057 001	10278936	Jun 28, 2033				
	10278937	Jun 28, 2033				
	10383840	Jun 28, 2033				
	10555924	Jun 28, 2033				
	10555925	Jun 28, 2033				
	8188146	Jan 27, 2020	DS DP			
	8293727	Feb 09, 2030			U-1287	
	8293728	Feb 09, 2030			U-1287	
	8298554	Apr 29, 2030	DP			
	8314086	Feb 09, 2030			U-1287	
	8318715	Feb 09, 2030			U-1287	
	8357677	Feb 09, 2030			U-1287	
	8367652	Feb 09, 2030			U-1287	
	8377920	Feb 09, 2030			U-1287	
	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	
	9918954	Jun 28, 2033			U-2699	
<u>ICOSAPENT ETHYL - VASCEPA</u>						
N 202057 002	10010517	Apr 29, 2030			I-819	Dec 13, 2022
	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	
	8188146	Jan 27, 2020	DS DP			
	8293727	Feb 09, 2030			U-1287	
	8293728	Feb 09, 2030			U-1287	
	8298554	Apr 29, 2030	DP			
	8314086	Feb 09, 2030			U-1287	
	8318715	Feb 09, 2030			U-1287	
	8357677	Feb 09, 2030			U-1287	

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<u>ICOSAPENT ETHYL - VASCEPA</u>						
N 202057	002	8367652	Feb 09, 2030	U-1287		
		8377920	Feb 09, 2030	U-1287		
		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		
		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	6800620	Apr 24, 2021	DS	U-1560	ODE-70 Jul 23, 2021
		6949535	Apr 24, 2021	DS	U-1560	ODE-71 Jul 23, 2021
		8138195	Apr 24, 2021	DS DP	U-1549	
		8492389	Apr 24, 2021	DS DP		
		8637533	Apr 24, 2021	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	6800620	Apr 24, 2021	DS	U-1560	ODE-70 Jul 23, 2021
		6949535	Apr 24, 2021	DS	U-1560	ODE-71 Jul 23, 2021
		8138195	Apr 24, 2021	DS DP	U-1549	
		8492389	Apr 24, 2021	DS DP		
		8637533	Apr 24, 2021	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		

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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 001	8586610	Nov 02, 2027	U-1625			
	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 002	8586610	Nov 02, 2027	U-1625			
	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			
	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			
	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			
	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			
	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 007	8586610	Nov 02, 2027	U-1625			
	8652776	Aug 31, 2030	U-1685			
	8999638	Oct 28, 2030	U-1674			
	9072742	Jan 16, 2031	U-1674			

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<u>ILOPERIDONE - FANAPT</u>						
N 022192	007	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>IMATINIB MESYLATE - GLEEVEC</u>						
N 021335	001	6958335	Dec 19, 2021	U-791		
<u>IMATINIB MESYLATE - GLEEVEC</u>						
N 021335	002	6958335	Dec 19, 2021	U-791		
<u>IMATINIB MESYLATE - GLEEVEC</u>						
N 021588	001	6958335	Dec 19, 2021	U-1883	ODE-40	Jan 25, 2020
		6958335	Dec 19, 2021	U-791		
		6958335*PED	Jun 19, 2022			
<u>IMATINIB MESYLATE - GLEEVEC</u>						
N 021588	002	6958335	Dec 19, 2021	U-1883	ODE-40	Jan 25, 2020
		6958335	Dec 19, 2021	U-791		
		6958335*PED	Jun 19, 2022			
<u>IMIQIUMOD - ALDARA</u>						
N 020723	001	7696159	Apr 01, 2024	DS U-1047		
		7696159	Apr 01, 2024	DS U-1048		
<u>IMIQIUMOD - ZYCLARA</u>						
N 022483	001	10238644	Dec 11, 2029	U-68		
		10238645	Aug 18, 2029	U-1455		
		10238645	Aug 18, 2029	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>IMIQIUMOD - ZYCLARA</u>						
N 022483	002	8222270	Dec 11, 2029	U-68		
<u>INDACATEROL MALEATE - ARCAPTA NEOHALER</u>						
N 022383	001	6878721	Feb 25, 2025	DS DP U-1168		
		8067437	Jun 02, 2020	U-1168		
		8479730	Oct 11, 2028	DP		
		8658673	Jun 02, 2020	DS DP U-1168		
		8796307	Jun 02, 2020	DS DP		
<u>INDINAVIR SULFATE - CRIXIVAN</u>						
N 020685	001	6689761	Feb 10, 2021	U-554		
<u>INDINAVIR SULFATE - CRIXIVAN</u>						
N 020685	003	6689761	Feb 10, 2021	U-554		
<u>INDINAVIR SULFATE - CRIXIVAN</u>						
N 020685	005	6689761	Feb 10, 2021	U-554		
<u>INDINAVIR SULFATE - CRIXIVAN</u>						
N 020685	006	6689761	Feb 10, 2021	U-554		
<u>INDOCYANINE GREEN - SPY AGENT GREEN KIT</u>						
N 211580	001	6915154	Aug 01, 2021	U-2460	NP	Nov 21, 2021
		7881777	Sep 22, 2021	U-2461		
		8185176	Jun 04, 2028	U-2462		
		8406860	Apr 09, 2029	U-2463		
		8647605	Feb 11, 2029	U-2464		
		8647605	Feb 11, 2029	U-2468		
		8892190	Aug 11, 2020	U-2465		
		9421280	Nov 24, 2025	U-2466		
		9421280	Nov 24, 2025	U-2467		

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<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>INGENOL MEBUTATE - PICATO</u>						
N 202833	001	7410656	Oct 10, 2020		U-1222	
		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	7410656	Oct 10, 2020		U-1222	
		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	7015315	Mar 21, 2023	DS		NCE Oct 05, 2023
		7101993	Sep 05, 2023	DS		ODE-212 Oct 05, 2025
		8101743	Apr 01, 2025	DS DP		
		8697860	Apr 29, 2031	DP		
		9061044	Apr 29, 2031	DS		
		9399774	Apr 29, 2031		U-2430	
<u>INSULIN ASPART - FIASP</u>						
N 208751	001	8324157	Jun 25, 2030	DP		M-247 Oct 21, 2022
						NP Sep 29, 2020
						NPP Dec 19, 2022
<u>INSULIN ASPART - FIASP FLEXTOUCH</u>						
N 208751	002	10220155	Jul 17, 2026	DP		NP Sep 29, 2020
		10357616	Jan 20, 2026	DP		NPP Dec 19, 2022
		10376652	Jan 20, 2026	DP		
		6899699	Jan 02, 2022	DP		
		7762994	May 23, 2024	DP		
		8324157	Jun 25, 2030	DP		
		8579869	Jun 30, 2023	DP		
		8672898	Jan 02, 2022	DP		
		8684969	Oct 20, 2025	DP		
		8920383	Jul 17, 2026	DP		
		9108002	Jan 20, 2026	DP		
		9132239	Feb 01, 2032	DP		
		9457154	Jan 20, 2026	DP		
		9486588	Jan 02, 2022	DP		
		9616180	Jan 20, 2026	DP		
		9687611	Feb 27, 2027	DP		
		9775953	Jul 17, 2026	DP		
		9861757	Jan 20, 2026	DP		

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APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>INSULIN ASPART - FIASP FLEXTOUCH</u>						
N 208751	002	RE46363	Aug 03, 2026	DP		
<u>INSULIN ASPART - FIASP PENFILL</u>						
N 208751	003				NPP	Dec 19, 2022
<u>INSULIN ASPART PROTAMINE RECOMBINANT; INSULIN ASPART RECOMBINANT - NOVOLOG MIX 70/30 FLEXPEN</u>						
N 021172	004	7762994	May 23, 2024	DP		
		8579869	Jun 30, 2023	DP		
		9265893	Sep 23, 2032	DP		
		RE41956	Jan 21, 2021	DP		
<u>INSULIN ASPART RECOMBINANT - NOVOLOG PENFILL</u>						
N 020986	002	7762994	May 23, 2024	DP		
		8579869	Jun 30, 2023	DP		
<u>INSULIN ASPART RECOMBINANT - NOVOLOG FLEXPEN</u>						
N 020986	003	7762994	May 23, 2024	DP		
		8579869	Jun 30, 2023	DP		
		9265893	Sep 23, 2032	DP		
		RE41956	Jan 21, 2021	DP		
<u>INSULIN ASPART RECOMBINANT - NOVOLOG INNOLET</u>						
N 020986	004	RE41956	Jan 21, 2021	DP		
<u>INSULIN ASPART RECOMBINANT - NOVOLOG FLEXTOUCH</u>						
N 020986	005	10220155	Jul 17, 2026	DP		
		10357616	Jan 20, 2026	DP		
		10376652	Jan 20, 2026	DP		
		6899699	Jan 02, 2022	DP		
		7762994	May 23, 2024	DP		
		8579869	Jun 30, 2023	DP		
		8672898	Jan 02, 2022	DP		
		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		
		9486588	Jan 02, 2022	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>INSULIN ASPART; INSULIN DEGLUDEC - RYZODEG 70/30</u>						
N 203313	001	10220155	Jul 17, 2026	DP	NCE	Sep 25, 2020
		10357616	Jan 20, 2026	DP		
		10376652	Jan 20, 2026	DP		
		6899699	Jan 02, 2022	DP		
		7615532	Jun 28, 2029	DS		
		7762994	May 23, 2024	DP		
		8579869	Jun 30, 2023	DP		
		8672898	Jan 02, 2022	DP		
		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		
		9486588	Jan 02, 2022	DP		
		9616180	Jan 20, 2026	DP		
		9687611	Feb 27, 2027	DP		
		9775953	Jul 17, 2026	DP		
		9861757	Jan 20, 2026	DP		
		9884094	May 01, 2033		U-2238	
		RE46363	Aug 03, 2026	DP		
<u>INSULIN DEGLUDEC - TRESIBA</u>						
N 203314	001	10220155	Jul 17, 2026	DP	NCE	Sep 25, 2020
		10357616	Jan 20, 2026	DP		
		10376652	Jan 20, 2026	DP		

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<u>INSULIN DEGLUDEC - TRESIBA</u>						
N 203314 001	6899699	Jan 02, 2022	DP			
	7615532	Jun 28, 2029	DS DP			
	7762994	May 23, 2024	DP			
	8579869	Jun 30, 2023	DP			
	8672898	Jan 02, 2022	DP			
	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			
	9486588	Jan 02, 2022	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>INSULIN DEGLUDEC - TRESIBA</u>						
N 203314 002	10220155	Jul 17, 2026	DP		NCE	Sep 25, 2020
	10357616	Jan 20, 2026	DP			
	10376652	Jan 20, 2026	DP			
	6899699	Jan 02, 2022	DP			
	7615532	Jun 28, 2029	DS DP			
	7762994	May 23, 2024	DP			
	8579869	Jun 30, 2023	DP			
	8672898	Jan 02, 2022	DP			
	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			
	9486588	Jan 02, 2022	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>INSULIN DEGLUDEC - TRESIBA</u>						
N 203314 003	10376652	Jan 20, 2026	DP			
	7615532	Jun 28, 2029	DS DP			
<u>INSULIN DEGLUDEC; LIRAGLUTIDE - XULTOPHY 100/3.6</u>						
N 208583 001	10220155	Jul 17, 2026	DP		M-242	Aug 08, 2022
	10357616	Jan 20, 2026	DP		NCE	Sep 25, 2020
	10376652	Jan 20, 2026	DP			
	6268343	Aug 22, 2022	DS DP			
	6268343*PED	Feb 22, 2023				
	6899699	Jan 02, 2022	DP			
	7615532	Jun 28, 2029	DS DP			
	7762994	May 23, 2024	DP			
	7762994*PED	Nov 23, 2024				
	8579869	Jun 30, 2023	DP			
	8579869*PED	Dec 30, 2023				
	8672898	Jan 02, 2022	DP			
	8684969	Oct 20, 2025	DP			
	8846618	Jun 27, 2022	DS DP			
	8846618*PED	Dec 27, 2022				
	8920383	Jul 17, 2026	DP			
	8937042	May 05, 2029	DP			
	9108002	Jan 20, 2026	DP			
	9132239	Feb 01, 2032	DP			
	9457154	Sep 27, 2027	DP			
	9486588	Jan 02, 2022	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			

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<u>INSULIN DETEMIR RECOMBINANT - LEVEMIR FLEXPEN</u>						
N 021536 002	9265893	Sep 23, 2032	DP			
	RE41956	Jan 21, 2021	DP			
<u>INSULIN DETEMIR RECOMBINANT - LEVEMIR FLEXTOUCH</u>						
N 021536 005	10220155	Jul 17, 2026	DP			
	10357616	Jan 20, 2026	DP			
	10376652	Jan 20, 2026	DP			
	6899699	Jan 02, 2022	DP			
	7686786	Aug 03, 2026	DP			
	7762994	May 23, 2024	DP			
	8579869	Jun 30, 2023	DP			
	8672898	Jan 02, 2022	DP			
	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			
	9486588	Jan 02, 2022	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>INSULIN GLARGINE RECOMBINANT - LANTUS</u>						
N 021081 001	7476652	Jul 23, 2023	DP			
	7713930	Jun 13, 2023	DP			
	7918833	Sep 23, 2027	DP			
<u>INSULIN GLARGINE RECOMBINANT - LANTUS SOLOSTAR</u>						
N 021081 002	8512297	Sep 15, 2024	DP			
	8556864	Mar 03, 2024	DP			
	8603044	Mar 02, 2024	DP			
	8679069	Apr 12, 2025	DP			
	8992486	Jun 05, 2024	DP			
	9011391	Mar 26, 2024			U-1832	
	9233211	Mar 02, 2024	DP			
	9408979	Mar 02, 2024	DP			
	9526844	Mar 02, 2024	DP			
	9533105	Aug 17, 2024	DP			
	9561331	Aug 28, 2024	DP			
	9604008	Mar 02, 2024	DP			
	9604009	Aug 16, 2024	DP			
	9610409	Mar 02, 2024	DP			
	9623189	Aug 19, 2024	DP			
	9717852	Apr 08, 2033	DP			
	9775954	Mar 02, 2024	DP			
	9827379	Mar 02, 2024	DP		U-2146	
<u>INSULIN GLARGINE RECOMBINANT - TOUJEO SOLOSTAR</u>						
N 206538 001	10369291	Sep 16, 2035	DP		NPP	Nov 26, 2022
	7918833	Sep 23, 2027	DP			
	7918833*PED	Mar 23, 2028				
	8512297	Sep 15, 2024	DP			
	8556864	Mar 03, 2024	DP			
	8603044	Mar 02, 2024	DP			
	8679069	Apr 12, 2025	DP			
	8992486	Jun 05, 2024	DP			
	9011391	Mar 26, 2024			U-1832	
	9233211	Mar 02, 2024	DP			
	9345750	May 18, 2031	DP		U-1855	
	9408979	Mar 02, 2024	DP			
	9526844	Mar 02, 2024	DP			
	9533105	Aug 17, 2024	DP			
	9561331	Aug 28, 2024	DP			
	9604008	Mar 02, 2024	DP			
	9604009	Aug 16, 2024	DP			
	9610409	Mar 02, 2024	DP			
	9623189	Aug 19, 2024	DP			
	9775954	Mar 02, 2024	DP			

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<u>INSULIN GLARGINE RECOMBINANT - TOUJEO SOLOSTAR</u>						
N 206538	001	9827379	Mar 02, 2024	DP		U-2146
<u>INSULIN GLARGINE RECOMBINANT - TOUJEO MAX SOLOSTAR</u>						
N 206538	002	10369291	Sep 16, 2035	DP	NPP	Nov 26, 2022
		7918833	Sep 23, 2027	DP		
		8512297	Sep 15, 2024	DP		
		8556864	Mar 03, 2024	DP		
		8603044	Mar 02, 2024	DP		
		8679069	Apr 12, 2025	DP		
		8992486	Jun 05, 2024	DP		
		9011391	Mar 26, 2024	DP		U-1832
		9233211	Mar 02, 2024	DP		
		9345750	May 18, 2031	DP		U-1855
		9408979	Mar 02, 2024	DP		
		9526844	Mar 02, 2024	DP		
		9533105	Aug 17, 2024	DP		
		9561331	Aug 28, 2024	DP		
		9604008	Mar 02, 2024	DP		
		9604009	Aug 16, 2024	DP		
		9610409	Mar 02, 2024	DP		
		9623189	Aug 19, 2024	DP		
		9775954	Mar 02, 2024	DP		
		9827379	Mar 02, 2024	DP		U-2146
<u>INSULIN GLARGINE; LIXISENATIDE - SOLIQUA 100/33</u>						
N 208673	001	10029011	Nov 11, 2030	DP	NCE	Jul 27, 2021
		10117909	Oct 09, 2029	DP		
		7918833	Sep 23, 2027	DP		
		8512297	Sep 15, 2024	DP		
		8556864	Mar 03, 2024	DP		
		8603044	Mar 02, 2024	DP		
		8679069	Apr 12, 2025	DP		
		8992486	Jun 05, 2024	DP		
		9011391	Mar 26, 2024			U-1923
		9011391	Mar 26, 2024			U-2496
		9233211	Mar 02, 2024	DP		
		9408979	Mar 02, 2024	DP		
		9526764	Oct 09, 2029	DP		
		9526844	Mar 02, 2024	DP		
		9533105	Aug 17, 2024	DP		
		9561331	Aug 28, 2024	DP		
		9604008	Mar 02, 2024	DP		
		9604009	Aug 16, 2024	DP		
		9610409	Mar 02, 2024	DP		
		9623189	Aug 19, 2024	DP		
		9707176	Nov 11, 2030	DP		
		9717852	Apr 08, 2033	DP		
		9775954	Mar 02, 2024	DP		
		9821032	May 09, 2032			U-2182
		9827379	Mar 02, 2024	DP		U-2146
		9950039	Dec 10, 2035			U-2277
		9950039	Dec 10, 2035			U-2278
		9950039	Dec 10, 2035			U-2279
		RE45313	Jul 12, 2020	DS DP		
<u>INSULIN GLULISINE RECOMBINANT - APIDRA</u>						
N 021629	001	6960561	Jan 25, 2023	DP		U-471
		7452860	Mar 22, 2022	DP		
		7696162	Mar 22, 2022	DP		U-471
<u>INSULIN GLULISINE RECOMBINANT - APIDRA</u>						
N 021629	002	6960561	Jan 25, 2023	DP		U-471
		7452860	Mar 22, 2022	DP		
		7696162	Mar 22, 2022	DP		U-471
<u>INSULIN GLULISINE RECOMBINANT - APIDRA SOLOSTAR</u>						
N 021629	003	6960561	Jan 25, 2023	DP		U-471
		7452860	Mar 22, 2022	DP		
		7696162	Mar 22, 2022	DP		U-471
		7918833	Sep 23, 2027	DP		

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APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>INSULIN GLULISINE RECOMBINANT - APIDRA SOLOSTAR</u>						
N 021629 003	8512297	Sep 15, 2024	DP			
	8556864	Mar 03, 2024	DP			
	8603044	Mar 02, 2024	DP			
	8679069	Apr 12, 2025	DP			
	8992486	Jun 05, 2024	DP			
	9011391	Mar 26, 2024	DP	U-1832		
	9233211	Mar 02, 2024	DP			
	9408979	Mar 02, 2024	DP			
	9526844	Mar 02, 2024	DP			
	9533105	Aug 17, 2024	DP			
	9561331	Aug 28, 2024	DP			
	9604008	Mar 02, 2024	DP			
	9604009	Aug 16, 2024	DP			
	9610409	Mar 02, 2024	DP			
	9623189	Aug 19, 2024	DP			
	9717852	Apr 08, 2033	DP			
	9775954	Mar 02, 2024	DP			
	9827379	Mar 02, 2024	DP	U-2146		
<u>INSULIN HUMAN - HUMULIN R</u>						
N 018780 004	7291132	Aug 09, 2024	DP			
<u>INSULIN LISPRO - ADMELOG</u>						
N 209196 001					NP	Dec 11, 2020
<u>INSULIN LISPRO - ADMELOG SOLOSTAR</u>						
N 209196 002	7918833	Sep 23, 2027	DP		NP	Dec 11, 2020
	8512297	Sep 15, 2024	DP			
	8556864	Mar 03, 2024	DP			
	8603044	Mar 02, 2024	DP			
	8679069	Apr 12, 2025	DP			
	8992486	Jun 05, 2024	DP			
	9011391	Mar 26, 2024	DP	U-1832		
	9233211	Mar 02, 2024	DP			
	9408979	Mar 02, 2024	DP			
	9526844	Mar 02, 2024	DP			
	9533105	Aug 17, 2024	DP			
	9561331	Aug 28, 2024	DP			
	9604008	Mar 02, 2024	DP			
	9604009	Aug 16, 2024	DP			
	9610409	Mar 02, 2024	DP			
	9623189	Aug 19, 2024	DP			
	9717852	Apr 08, 2033	DP			
	9775954	Mar 02, 2024	DP			
	9827379	Mar 04, 2024	DP	U-2146		
<u>INSULIN LISPRO PROTAMINE RECOMBINANT; INSULIN LISPRO RECOMBINANT - HUMALOG MIX 75/25 KWIKPEN</u>						
N 021017 002	7291132	Aug 09, 2024	DP			
<u>INSULIN LISPRO PROTAMINE RECOMBINANT; INSULIN LISPRO RECOMBINANT - HUMALOG MIX 50/50 KWIKPEN</u>						
N 021018 002	7291132	Aug 09, 2024	DP			
<u>INSULIN LISPRO RECOMBINANT - HUMALOG KWIKPEN</u>						
N 020563 003	7291132	Aug 09, 2024	DP			
<u>INSULIN LISPRO RECOMBINANT - HUMALOG KWIKPEN</u>						
N 205747 001	7291132	Aug 09, 2024	DP			
<u>INSULIN RECOMBINANT HUMAN - EXUBERA</u>						
N 021868 001	6582728	Jun 24, 2020	DP			
<u>INSULIN RECOMBINANT HUMAN - EXUBERA</u>						
N 021868 002	6582728	Jun 24, 2020	DP			
<u>INSULIN RECOMBINANT HUMAN - AFREZZA</u>						
N 022472 001	10046031	Aug 11, 2029		U-2383		
	10201672	Aug 02, 2030	DP			
	10342938	Jun 12, 2029	DP			
	10500159	Nov 02, 2030	DP			
	6444226	Jun 29, 2020	DP	U-1534		

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<u>INSULIN RECOMBINANT HUMAN - AFREZZA</u>						
N 022472 001	6652885	Jun 29, 2020				U-1535
	7305986	Jan 16, 2023	DP			
	7464706	Mar 02, 2023	DP			
	7648960	Jun 29, 2020			U-1535	
	7943178	Jun 29, 2020	DP		U-1535	
	7943572	Aug 10, 2026			U-1539	
	8119593	Aug 11, 2029			U-1537	
	8146588	Apr 24, 2023	DP			
	8156936	Jan 16, 2023	DP			
	8215300	Nov 24, 2022	DP			
	8258095	Aug 11, 2029			U-1537	
	8389470	Jun 29, 2020	DP		U-1621	
	8424518	Oct 17, 2031	DP			
	8485180	Mar 25, 2030	DP			
	8499757	Feb 19, 2032	DP			
	8551528	Jun 11, 2030	DP			
	8623817	Sep 18, 2029			U-1537	
	8636001	Jul 12, 2032	DP			
	8729019	Dec 26, 2028	DP			
	8734845	Jun 11, 2030	DP			
	8778403	Jun 11, 2030	DP		U-1538	
	8889099	Jun 29, 2020	DP		U-1621	
	8912193	Jun 12, 2029	DP		U-1538	
	8950397	Jul 20, 2021	DP			
	9192675	Jun 12, 2029	DP		U-1788	
	9283193	Sep 14, 2026	DP			
	9339615	Oct 20, 2029	DP			
	9358352	Feb 15, 2031	DP		U-1861	
	9393372	Jul 04, 2029	DP			
	9446133	Jun 12, 2029	DP		U-1861	
	9511198	Feb 16, 2030			U-1929	
	9511198	Feb 16, 2030			U-1930	
	9597374	Oct 08, 2031			U-1987	
	9662461	Jun 12, 2029	DP		U-2019	
	9717689	Sep 14, 2026	DP			
	9943571	Aug 11, 2029			U-1537	
<u>INSULIN RECOMBINANT HUMAN - AFREZZA</u>						
N 022472 002	10046031	Aug 11, 2029				U-2383
	10201672	Aug 02, 2030	DP			
	10342938	Jun 12, 2029	DP			
	10500159	Nov 02, 2030	DP			
	6444226	Jun 29, 2020	DP		U-1534	
	6652885	Jun 29, 2020			U-1535	
	7305986	Jan 16, 2023	DP			
	7464706	Mar 02, 2023	DP			
	7648960	Jun 29, 2020			U-1535	
	7943178	Jun 29, 2020	DP		U-1535	
	7943572	Aug 10, 2026			U-1539	
	8119593	Aug 11, 2029			U-1537	
	8146588	Apr 24, 2023	DP			
	8156936	Jan 16, 2023	DP			
	8215300	Nov 24, 2022	DP			
	8258095	Aug 11, 2029			U-1537	
	8389470	Jun 29, 2020	DP		U-1621	
	8424518	Oct 17, 2031	DP			
	8485180	Mar 25, 2030	DP			
	8499757	Feb 19, 2032	DP			
	8551528	Jun 11, 2030	DP			
	8623817	Sep 18, 2029			U-1537	
	8636001	Jul 12, 2032	DP			
	8729019	Dec 26, 2028	DP			
	8734845	Jun 11, 2030	DP			
	8778403	Jun 11, 2030	DP		U-1538	
	8889099	Jun 29, 2020	DP		U-1621	
	8912193	Jun 12, 2029	DP		U-1538	
	8950397	Jul 20, 2021	DP			
	9192675	Jun 12, 2029	DP		U-1788	
	9283193	Sep 14, 2026	DP			

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<u>INSULIN RECOMBINANT HUMAN - AFREZZA</u>						
N 022472 002	9339615	Oct 20, 2029	DP			
	9358352	Feb 15, 2031	DP U-1861			
	9393372	Jul 04, 2029	DP			
	9446133	Jun 12, 2029	DP U-1861			
	9511198	Feb 16, 2030		U-1929		
	9511198	Feb 16, 2030		U-1930		
	9597374	Oct 08, 2031		U-1987		
	9662461	Jun 12, 2029	DP U-2019			
	9717689	Sep 14, 2026	DP			
	9943571	Aug 11, 2029		U-1537		
<u>INSULIN RECOMBINANT HUMAN - AFREZZA</u>						
N 022472 003	10046031	Aug 11, 2029		U-2383		
	10201672	Aug 02, 2030	DP			
	10342938	Jun 12, 2029	DP			
	10500159	Nov 02, 2030	DP			
	6444226	Jun 29, 2020	DP U-1534			
	6652885	Jun 29, 2020		U-1535		
	7305986	Jan 16, 2023	DP			
	7464706	Mar 02, 2023	DP			
	7648960	Jun 29, 2020		U-1535		
	7943178	Jun 29, 2020	DP U-1535			
	7943572	Aug 10, 2026		U-1539		
	8119593	Aug 11, 2029		U-1537		
	8146588	Apr 24, 2023	DP			
	8156936	Jan 16, 2023	DP			
	8215300	Nov 24, 2022	DP			
	8258095	Aug 11, 2029		U-1537		
	8389470	Jun 29, 2020	DP U-1621			
	8424518	Oct 17, 2031	DP			
	8485180	Mar 25, 2030	DP			
	8499757	Feb 19, 2032	DP			
	8551528	Jun 11, 2030	DP			
	8623817	Sep 18, 2029		U-1537		
	8636001	Jul 12, 2032	DP			
	8729019	Dec 26, 2028	DP			
	8734845	Jun 11, 2030	DP			
	8778403	Jun 11, 2030	DP U-1538			
	8889099	Jun 29, 2020	DP U-1621			
	8912193	Jun 12, 2029	DP U-1538			
	8950397	Jul 20, 2021	DP			
	9192675	Jun 12, 2029	DP U-1788			
	9283193	Sep 14, 2026	DP			
	9339615	Oct 20, 2029	DP			
	9358352	Feb 15, 2031	DP U-1861			
	9393372	Jul 04, 2029	DP			
	9446133	Jun 12, 2029	DP U-1861			
	9511198	Feb 16, 2030		U-1929		
	9511198	Feb 16, 2030		U-1930		
	9597374	Oct 08, 2031		U-1987		
	9662461	Jun 12, 2029	DP U-2019			
	9717689	Sep 14, 2026	DP			
	9943571	Aug 11, 2029		U-1537		
<u>INSULIN RECOMBINANT HUMAN; INSULIN SUSP ISOPHANE RECOMBINANT HUMAN - HUMULIN 70/30</u>						
N 019717 001	7291132	Aug 09, 2024	DP			
<u>INSULIN RECOMBINANT HUMAN; INSULIN SUSP ISOPHANE RECOMBINANT HUMAN - HUMULIN 70/30 PEN</u>						
N 019717 002	7291132	Aug 09, 2024	DP			
<u>INSULIN SUSP ISOPHANE RECOMBINANT HUMAN - HUMULIN N</u>						
N 018781 001	7291132	Aug 09, 2024	DP			
<u>IOBENGUANE I-131 - AZEDRA</u>						
N 209607 001				NP		Jul 30, 2021
				ODE-204		Jul 30, 2025

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<u>IODIXANOL - VISIPAQUE 320</u>						
N 020351	002				I-752	Apr 05, 2020
<u>IODIXANOL - VISIPAQUE 320</u>						
N 020808	002				I-752	Apr 05, 2020
<u>IPRATROPIUM BROMIDE - ATROVENT HFA</u>						
N 021527	001	6739333	May 26, 2020	DP		
		6983743	May 26, 2020	DP		
		8474447	Jan 17, 2030	DP		
<u>IRINOTECAN HYDROCHLORIDE - CAMPTOSAR</u>						
N 020571	001	6403569	Apr 28, 2020	U-449		
		6794370	May 01, 2020	U-606		
<u>IRINOTECAN HYDROCHLORIDE - CAMPTOSAR</u>						
N 020571	002	6403569	Apr 28, 2020	U-449		
		6794370	May 01, 2020	U-606		
<u>IRINOTECAN HYDROCHLORIDE - ONIVYDE</u>						
N 207793	001	10456360	Oct 15, 2036	DP	ODE-99	Oct 22, 2022
		8147867	Aug 29, 2028	DS DP		
		8329213	May 02, 2025	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	6812238	Oct 31, 2020	DS	NCE	Mar 06, 2020
		7459561	Oct 31, 2020	DS	ODE-90	Mar 06, 2022
					GAIN	Mar 06, 2025
					GAIN	Mar 06, 2027
<u>ISAVUCONAZONIUM SULFATE - CRESEMBA</u>						
N 207501	001	6812238	Oct 31, 2020	DS	NCE	Mar 06, 2020
		7459561	Oct 31, 2020	DS	ODE-90	Mar 06, 2022
					GAIN	Mar 06, 2025
					GAIN	Mar 06, 2027
<u>ISOPROPYL ALCOHOL - ZURAGARD</u>						
N 210872	001				NP	Apr 26, 2022
<u>ISOTRETINOIN - ABSORICA</u>						
N 021951	001	7435427	Sep 21, 2021	DP		
		8367102	Sep 21, 2021		U-1347	
		8952064	Sep 21, 2021	DP		
		9078925	Sep 21, 2021	DP		
		9089534	Sep 21, 2021	DP		
<u>ISOTRETINOIN - ABSORICA</u>						
N 021951	002	7435427	Sep 21, 2021	DP		
		8367102	Sep 21, 2021		U-1347	
		8952064	Sep 21, 2021	DP		
		9078925	Sep 21, 2021	DP		
		9089534	Sep 21, 2021	DP		
<u>ISOTRETINOIN - ABSORICA</u>						
N 021951	003	7435427	Sep 21, 2021	DP		
		8367102	Sep 21, 2021		U-1347	
		8952064	Sep 21, 2021	DP		
		9078925	Sep 21, 2021	DP		

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<u>ISOTRETINOIN - ABSORICA</u>						
N 021951	003 9089534	Sep 21, 2021	DP			
<u>ISOTRETINOIN - ABSORICA</u>						
N 021951	004 7435427	Sep 21, 2021	DP			
	8367102	Sep 21, 2021	DP	U-1347		
	8952064	Sep 21, 2021	DP			
	9078925	Sep 21, 2021	DP			
	9089534	Sep 21, 2021	DP			
<u>ISOTRETINOIN - ABSORICA</u>						
N 021951	005 7435427	Sep 21, 2021	DP			
	8367102	Sep 21, 2021	DP	U-1347		
	8952064	Sep 21, 2021	DP			
	9078925	Sep 21, 2021	DP			
	9089534	Sep 21, 2021	DP			
<u>ISOTRETINOIN - ABSORICA</u>						
N 021951	006 7435427	Sep 21, 2021	DP			
	8367102	Sep 21, 2021	DP	U-1347		
	8952064	Sep 21, 2021	DP			
	9078925	Sep 21, 2021	DP			
	9089534	Sep 21, 2021	DP			
<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			
<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 7541363	Nov 13, 2024	DS DP		NCE	Aug 27, 2024
	7727993	Jan 28, 2023		U-2623		
	7727994	Jan 18, 2023		U-2623		
	8318201	Sep 05, 2027	DP			
<u>ISTRADEFYLLINE - NOURIANZ</u>						
N 022075	002 7541363	Nov 13, 2024	DS DP		NCE	Aug 27, 2024
	7727993	Jan 28, 2023		U-2623		
	7727994	Jan 18, 2023		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		
	8771739	Jul 25, 2023	DP			
	8921374	Jun 21, 2033	DP			
	9272046	Jun 21, 2033	DP			
	9713642	Jun 21, 2033		U-2453		

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<u>IVABRADINE - CORLANOR</u>						
N 209964 001	7361649	Feb 22, 2026	DS DP U-1694		NCE	Apr 15, 2020
	7361650	Feb 22, 2026	DS DP U-1694		NP	Apr 22, 2022
	7867996	Feb 22, 2026	DS DP U-1694		ODE-234	Apr 22, 2026
	7879842	Feb 22, 2026	DS DP U-1694		PED	Oct 15, 2020
					PED	Oct 22, 2022
					PED	Oct 22, 2026
<u>IVABRADINE HYDROCHLORIDE - CORLANOR</u>						
N 206143 001	7361649	Feb 22, 2026	DS DP U-1694		NCE	Apr 15, 2020
	7361649*PED	Aug 22, 2026			PED	Oct 15, 2020
	7361650	Feb 22, 2026	DS DP U-1694			
	7361650*PED	Aug 22, 2026				
	7867996	Feb 22, 2026	DS DP U-1694			
	7867996*PED	Aug 22, 2026				
	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		NCE	Apr 15, 2020
	7361649*PED	Aug 22, 2026			PED	Oct 15, 2020
	7361650	Feb 22, 2026	DS DP U-1694			
	7361650*PED	Aug 22, 2026				
	7867996	Feb 22, 2026	DS DP U-1694			
	7867996*PED	Aug 22, 2026				
	7879842	Feb 22, 2026	DS DP U-1694			
	7879842*PED	Aug 22, 2026				
<u>IVACAFTOR - KALYDECO</u>						
N 203188 001	7495103	May 20, 2027	DS DP		NPP	Jul 31, 2020
	8324242	Aug 05, 2027		U-1311	ODE-186	Feb 21, 2021
	8324242	Aug 05, 2027		U-1906	ODE-187	Dec 29, 2021
	8354427	Jul 06, 2026		U-1311	ODE-189	Jul 31, 2024
	8354427	Jul 06, 2026		U-1905	ODE-190	May 17, 2024
	8410274	Dec 28, 2026		DP	ODE-199	Aug 15, 2025
	8629162	Jun 24, 2025		U-2234		
	8754224	Dec 28, 2026	DS DP			
	9670163	Dec 28, 2026	DP	U-1311		
<u>IVACAFTOR - KALYDECO</u>						
N 203188 002					NPP	Jul 31, 2020
					ODE-186	Feb 21, 2021
					ODE-187	Dec 29, 2021
					ODE-189	Jul 31, 2024
					ODE-190	May 17, 2024
					ODE-199	Aug 15, 2025
<u>IVACAFTOR - KALYDECO</u>						
N 207925 001	10272046	Feb 27, 2033	DP	U-2531	NPP	Jul 31, 2020
	7495103	May 20, 2027	DS DP		NPP	Apr 29, 2022
	8324242	Aug 05, 2027		U-1311	ODE-188	Mar 17, 2022
	8324242	Aug 05, 2027		U-1906	ODE-189	Jul 31, 2024
	8324242	Aug 05, 2027		U-2527	ODE-190	May 17, 2024
	8354427	Jul 06, 2026		U-1311	ODE-199	Aug 15, 2025
	8354427	Jul 06, 2026		U-1905	ODE-236	Apr 29, 2026
	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	NPP	Jul 31, 2020
	7495103	May 20, 2027	DS DP		NPP	Apr 29, 2022
	8324242	Aug 05, 2027		U-1311	ODE-188	Mar 17, 2022
	8324242	Aug 05, 2027		U-1906	ODE-189	Jul 31, 2024
	8324242	Aug 05, 2027		U-2527	ODE-190	May 17, 2024
	8354427	Jul 06, 2026		U-1311	ODE-199	Aug 15, 2025

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<u>IVACAFTOR - KALYDECO</u>						
N 207925 002	8354427	Jul 06, 2026		U-1905	ODE-236	Apr 29, 2026
	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 003	10272046	Feb 27, 2033	DP	U-2531	NPP	Jul 31, 2020
	7495103	May 20, 2027	DS DP		NPP	Apr 29, 2022
	8324242	Aug 05, 2027		U-1311	ODE-188	Mar 17, 2022
	8324242	Aug 05, 2027		U-1906	ODE-189	Jul 31, 2024
	8324242	Aug 05, 2027		U-2527	ODE-190	May 17, 2024
	8354427	Jul 06, 2026		U-1311	ODE-199	Aug 15, 2025
	8354427	Jul 06, 2026		U-1905	ODE-236	Apr 29, 2026
	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; IVACAFTOR, TEZACAFTOR - SYMDEKO (COPACKAGED)</u>						
N 210491 001	10022352	Apr 09, 2027	DP	U-2343	NCE	Feb 12, 2023
	10022352	Apr 09, 2027	DP	U-2573	NPP	Jun 21, 2022
	10058546	Jul 15, 2033		U-2399	ODE-173	Feb 12, 2025
	10058546	Jul 15, 2033		U-2572	ODE-247	Jun 21, 2026
	10081621	Mar 25, 2031	DP	U-2420		
	10081621	Mar 25, 2031	DP	U-2571		
	10206877	Apr 14, 2035	DP	U-2498		
	10206877	Apr 14, 2035	DP	U-2570		
	10239867	Apr 09, 2027	DS DP	U-2512		
	10239867	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		
	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	NCE	Feb 12, 2023
	10022352	Apr 09, 2027	DP	U-2573	NPP	Jun 21, 2022
	10058546	Jul 15, 2033		U-2399	ODE-173	Feb 12, 2025
	10058546	Jul 15, 2033		U-2572	ODE-247	Jun 21, 2026
	10081621	Mar 25, 2031	DP	U-2420		
	10081621	Mar 25, 2031	DP	U-2571		
	10206877	Apr 14, 2035	DP	U-2498		
	10206877	Apr 14, 2035	DP	U-2570		
	10239867	Apr 09, 2027	DS DP	U-2512		
	10239867	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		

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<u>IVACAFTOR; IVACAFTOR, TEZACAFTOR - SYMDEKO (COPACKAGED)</u>						
N 210491 002	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	M-218	Jan 25, 2021
	7495103	May 20, 2027	DS DP		NCE	Jul 02, 2020
	7973038	Nov 08, 2026		U-1973	ODE-123	Sep 28, 2023
	8324242	Aug 05, 2027		U-1311	ODE-93	Jul 02, 2022
	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	Dec 04, 2028		U-1717		
	8846718	Dec 04, 2028		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	7495103	May 20, 2027	DS DP		M-218	Jan 25, 2021
	7973038	Nov 08, 2026		U-1973	NCE	Jul 02, 2020
	8324242	Aug 05, 2027		U-1911	ODE-123	Sep 28, 2023
	8410274	Dec 28, 2026	DP		ODE-93	Jul 02, 2022
	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	Dec 04, 2028		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	7495103	May 20, 2027	DS DP		NCE	Jul 02, 2020
	7973038	Nov 08, 2026		U-2374	NP	Aug 07, 2021
	8324242	Aug 05, 2027		U-2374	ODE-195	Aug 07, 2025
	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	Dec 04, 2028		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		

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<u>IVACAFTOR; LUMACAFTOR - ORKAMBI</u>						
N 211358	001	9931334	Dec 28, 2026	DP		U-2376
<u>IVACAFTOR; LUMACAFTOR - ORKAMBI</u>						
N 211358	002	7495103	May 20, 2027	DS DP	NCE	Jul 02, 2020
		7973038	Nov 08, 2026		NP	Aug 07, 2021
		8324242	Aug 05, 2027		ODE-195	Aug 07, 2025
		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	Dec 04, 2028		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>IVERMECTIN - IVERMECTIN</u>						
A 210019	001				PC	Apr 11, 2020
<u>IVERMECTIN - SKLICE</u>						
N 202736	001	8791153	Oct 12, 2027	DP		
		8927595	Oct 12, 2027		U-1782	
<u>IVERMECTIN - SOOLANTRA</u>						
N 206255	001	10206939	Mar 13, 2034		U-1631	
		7550440	Apr 22, 2024	DP	U-1631	
		8080530	Apr 22, 2024	DP	U-1631	
		8093219	Apr 22, 2024	DP	U-1631	
		8415311	Apr 22, 2024	DP	U-1631	
		8470788	Apr 22, 2024	DP	U-1631	
		8815816	Apr 22, 2024	DP	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-816	May 02, 2022
		9474779	Aug 19, 2033	DS DP	U-2350	NCE Jul 20, 2023
		9474779	Aug 19, 2033	DS DP	U-2533	ODE-203 Jul 20, 2025
		9474779	Aug 19, 2033	DS DP	U-2534	ODE-242 May 02, 2026
		9850277	Jan 18, 2033	DS DP	U-2350	
		9850277	Jan 18, 2033	DS DP	U-2533	
		9850277	Jan 18, 2033	DS DP	U-2534	
		9968595	Mar 13, 2035	DP	U-2351	
		9968595	Mar 13, 2035	DP	U-2533	
		9968595	Mar 13, 2035	DP	U-2534	
<u>IXABEPILONE - IXEMPRA KIT</u>						
N 022065	001	6670384	Jan 23, 2022	DP	U-959	
		6670384	Jan 23, 2022	DP	U-960	
		7022330	Jan 23, 2022	DP	U-958	
		7312237	Aug 21, 2024		U-965	
		RE41393	Feb 08, 2022		U-961	
		RE41911	Sep 28, 2020	DS DP	U-961	
<u>IXABEPILONE - IXEMPRA KIT</u>						
N 022065	002	6670384	Jan 23, 2022	DP	U-959	
		6670384	Jan 23, 2022	DP	U-960	
		7022330	Jan 23, 2022	DP	U-958	
		7312237	Aug 21, 2024		U-965	
		RE41393	Feb 08, 2022		U-961	
		RE41911	Sep 28, 2020	DS DP	U-961	

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<u>IXAZOMIB CITRATE - NINLARO</u>						
N 208462 001	7442830	Nov 20, 2029	DS DP U-2434		NCE	Nov 20, 2020
	7687662	Aug 06, 2027	DS DP		ODE-103	Nov 20, 2022
	8003819	Aug 06, 2027	DS DP U-2434			
	8530694	Aug 06, 2027	DS DP U-2434			
	8546608	Aug 12, 2024	DS			
	8859504	Jun 16, 2029	DS DP			
	8871745	Aug 06, 2027	U-2434			
	9175017	Jun 16, 2029	U-2434			
	9233115	Aug 12, 2024	U-2434			
<u>IXAZOMIB CITRATE - NINLARO</u>						
N 208462 002	7442830	Nov 20, 2029	DS DP U-2434		NCE	Nov 20, 2020
	7687662	Aug 06, 2027	DS DP		ODE-103	Nov 20, 2022
	8003819	Aug 06, 2027	DS DP U-2434			
	8530694	Aug 06, 2027	DS DP U-2434			
	8546608	Aug 12, 2024	DS			
	8859504	Jun 16, 2029	DS DP			
	8871745	Aug 06, 2027	U-2434			
	9175017	Jun 16, 2029	U-2434			
	9233115	Aug 12, 2024	U-2434			
<u>IXAZOMIB CITRATE - NINLARO</u>						
N 208462 003	7442830	Nov 20, 2029	DS DP U-2434		NCE	Nov 20, 2020
	7687662	Aug 06, 2027	DS DP		ODE-103	Nov 20, 2022
	8003819	Aug 06, 2027	DS DP U-2434			
	8530694	Aug 06, 2027	DS DP U-2434			
	8546608	Aug 12, 2024	DS			
	8859504	Jun 16, 2029	DS DP			
	8871745	Aug 06, 2027	U-2434			
	9175017	Jun 16, 2029	U-2434			
	9233115	Aug 12, 2024	U-2434			
<u>KETOCONAZOLE - XOLEGEL</u>						
N 021946 001	8232276	Nov 24, 2020	DP			
<u>KETOROLAC TROMETHAMINE - ACULAR LS</u>						
N 021528 001	8008338	May 24, 2027	DS DP U-1181			
	8207215	May 28, 2024	U-1251			
	8377982	May 28, 2024	U-1363			
	8377982*PED	Nov 28, 2024				
	8541463	May 28, 2024	U-1441			
	8541463*PED	Nov 28, 2024				
	8648107	May 28, 2024	DP			
	8906950	May 28, 2024	U-1626			
	8946281	May 28, 2024	U-1662			
	9216167	May 28, 2024	U-1800			
<u>KETOROLAC TROMETHAMINE - ACUVAIL</u>						
N 022427 001	7842714	Aug 15, 2029	DS DP			
	8512717	Mar 07, 2028	DP			
	8992952	Aug 05, 2024	DP			
	9192571	Mar 07, 2028	DP			
<u>KETOROLAC TROMETHAMINE; PHENYLEPHRINE HYDROCHLORIDE - OMIIDRIA</u>						
N 205388 001	8173707	Jul 30, 2023	U-1518		NPP	Dec 08, 2020
	8173707*PED	Jan 30, 2024			PED	Jun 08, 2021
	8586633	Jul 30, 2023	DP			
	8586633*PED	Jan 30, 2024				
	9066856	Oct 23, 2033	DP			
	9066856*PED	Apr 23, 2034				
	9278101	Jul 30, 2023	U-1518			
	9278101*PED	Jan 30, 2024				
	9399040	Jul 30, 2023	DP			
	9399040*PED	Jan 30, 2024				
	9486406	Oct 23, 2033	DP			
	9486406*PED	Apr 23, 2034				
	9855246	Oct 23, 2033	DP			

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<u>L-GLUTAMINE - ENDARI</u>						
N 208587	001				I-748 ODE-150	Jul 07, 2020 Jul 07, 2024
<u>LACOSAMIDE - VIMPAT</u>						
N 022253	001	RE38551	Mar 17, 2022	DS DP U-1567	NPP	Nov 03, 2020
		RE38551	Mar 17, 2022	DS DP U-2140		
<u>LACOSAMIDE - VIMPAT</u>						
N 022253	002	RE38551	Mar 17, 2022	DS DP U-1567	NPP	Nov 03, 2020
		RE38551	Mar 17, 2022	DS DP U-2140		
<u>LACOSAMIDE - VIMPAT</u>						
N 022253	003	RE38551	Mar 17, 2022	DS DP U-1567	NPP	Nov 03, 2020
		RE38551	Mar 17, 2022	DS DP U-2140		
<u>LACOSAMIDE - VIMPAT</u>						
N 022253	004	RE38551	Mar 17, 2022	DS DP U-1567	NPP	Nov 03, 2020
		RE38551	Mar 17, 2022	DS DP U-2140		
<u>LACOSAMIDE - VIMPAT</u>						
N 022254	001	RE38551	Mar 17, 2022	DS DP U-1565	M-217	Nov 03, 2020
		RE38551	Mar 17, 2022	DS DP U-1568		
<u>LACOSAMIDE - VIMPAT</u>						
N 022255	001	RE38551	Mar 17, 2022	DS DP U-1567	NPP	Nov 03, 2020
		RE38551	Mar 17, 2022	DS DP U-2140		
<u>LACTITOL - PIZENSY</u>						
N 211281	001				NCE	Feb 12, 2025
<u>LAMIVUDINE; RALTEGRAVIR POTASSIUM - DUTREBIS</u>						
N 206510	001	7169780	Oct 03, 2023	DS DP		
		7169780*PED	Apr 03, 2024			
		7217713	Oct 21, 2022		U-1663	
		7217713*PED	Apr 21, 2023			
		7435734	Oct 21, 2022		U-1663	
		7435734*PED	Apr 21, 2023			
		7754731	Mar 11, 2029	DS DP	U-1663	
		7754731*PED	Sep 11, 2029			
		7820660	Apr 25, 2023	DS		
<u>LAMOTRIGINE - LAMICTAL XR</u>						
N 022115	001	8637512	Jun 14, 2028	DP		
		9144547	Sep 22, 2023	DP		
<u>LAMOTRIGINE - LAMICTAL XR</u>						
N 022115	002	8637512	Jun 14, 2028	DP		
		9144547	Sep 22, 2023	DP		
<u>LAMOTRIGINE - LAMICTAL XR</u>						
N 022115	003	8637512	Jun 14, 2028	DP		
		9144547	Sep 22, 2023	DP		
<u>LAMOTRIGINE - LAMICTAL XR</u>						
N 022115	004	8637512	Jun 14, 2028	DP		
		9144547	Sep 22, 2023	DP		
<u>LAMOTRIGINE - LAMICTAL XR</u>						
N 022115	005	8637512	Jun 14, 2028	DP		
		9144547	Sep 22, 2023	DP		
<u>LAMOTRIGINE - LAMICTAL XR</u>						
N 022115	006	8637512	Jun 14, 2028	DP		
		9144547	Sep 22, 2023	DP		
<u>LAMOTRIGINE - LAMICTAL ODT</u>						
N 022251	001	7919115	Jan 04, 2029	DS DP		
		8840925	Jul 02, 2028	DP	U-1596	
		9339504	Jul 02, 2028	DP	U-1596	

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<u>LAMOTRIGINE - LAMICTAL ODT</u>						
N 022251	004	7919115	Jan 04, 2029	DS DP		
		8840925	Jul 02, 2028	DP	U-1596	
		9339504	Jul 02, 2028	DP	U-1596	
<u>LANREOTIDE ACETATE - SOMATULINE DEPOT</u>						
N 022074	001	5595760	Mar 08, 2020	DP	U-831	I-754 Sep 15, 2020 ODE-156 Sep 15, 2024 ODE-82 Dec 16, 2021
<u>LANREOTIDE ACETATE - SOMATULINE DEPOT</u>						
N 022074	002	5595760	Mar 08, 2020	DP	U-831	I-754 Sep 15, 2020 ODE-156 Sep 15, 2024 ODE-82 Dec 16, 2021
<u>LANREOTIDE ACETATE - SOMATULINE DEPOT</u>						
N 022074	003	5595760	Mar 08, 2020	DP	U-831	I-754 Sep 15, 2020 ODE-156 Sep 15, 2024 ODE-82 Dec 16, 2021
<u>LANSOPRAZOLE - PREVACID IV</u>						
N 021566	001	7396841	Aug 17, 2021	DP	U-947	
<u>LANTHANUM CARBONATE - FOSRENOL</u>						
N 021468	001	7381428	Aug 26, 2024		U-890	
		7465465	Aug 26, 2024	DP		
<u>LANTHANUM CARBONATE - FOSRENOL</u>						
N 021468	002	7381428	Aug 26, 2024		U-890	
		7465465	Aug 26, 2024	DP		
<u>LANTHANUM CARBONATE - FOSRENOL</u>						
N 021468	003	7381428	Aug 26, 2024		U-890	
		7465465	Aug 26, 2024	DP		
<u>LANTHANUM CARBONATE - FOSRENOL</u>						
N 021468	004	7381428	Aug 26, 2024		U-890	
		7465465	Aug 26, 2024	DP		
<u>LANTHANUM CARBONATE - FOSRENOL</u>						
N 204734	001	7465465	Aug 26, 2024	DP		
		8980327	Dec 01, 2030	DP		
		9023397	Dec 01, 2030	DP		
<u>LANTHANUM CARBONATE - FOSRENOL</u>						
N 204734	002	7465465	Aug 26, 2024	DP		
		8980327	Dec 01, 2030	DP		
		9023397	Dec 01, 2030	DP		
<u>LAPATINIB DITOSYLATE - TYKERB</u>						
N 022059	001	6713485	Sep 29, 2020	DS DP	U-1429	M-235 Dec 06, 2021
		6713485	Sep 29, 2020	DS DP	U-800	
		7157466	Nov 19, 2021	DS DP		
		8821927	Sep 18, 2029	DS DP		
<u>LAROTRECTINIB SULFATE - VITRAKVI</u>						
N 210861	001	10005783	Oct 21, 2029		U-2472	NCE Nov 26, 2023
		10047097	Oct 21, 2029		U-2474	ODE-215 Nov 26, 2025
		10172861	Nov 16, 2035	DS DP		ODE-220 Nov 26, 2025
		10285993	Nov 16, 2035		U-2470	ODE-221 Nov 26, 2025
		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 210861	002	10005783	Oct 21, 2029		U-2472	NCE Nov 26, 2023
		10047097	Oct 21, 2029		U-2474	ODE-215 Nov 26, 2025
		10172861	Nov 16, 2035	DS DP		ODE-220 Nov 26, 2025
		10285993	Nov 16, 2035		U-2470	ODE-221 Nov 26, 2025

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<u>LAROTRECTINIB SULFATE - VITRAKVI</u>						
N 210861 002	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	NCE	Nov 26, 2023
	10045991	Apr 04, 2037		U-2473	ODE-215	Nov 26, 2025
	10047097	Oct 21, 2029		U-2474	ODE-220	Nov 26, 2025
	10137127	Apr 04, 2037	DP		ODE-221	Nov 26, 2025
	10172861	Nov 16, 2035	DS			
	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					NCE	Jan 31, 2025
<u>LASMIDITAN SUCCINATE - REYVOW</u>						
N 211280 002					NCE	Jan 31, 2025
<u>LATANOPROST - XELPROS</u>						
N 206185 001	9539262	Oct 15, 2028		U-2400		
	9629852	Sep 12, 2029	DP			
<u>LATANOPROST; NETARSUDIL DIMESYLATE - ROCKLATAN</u>						
N 208259 001	10174017	Jan 27, 2030	DS DP	U-1524	NC	Mar 12, 2022
	8394826	Nov 10, 2030	DS DP	U-1524	NCE	Dec 18, 2022
	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	Jan 05, 2025	DS	U-2144		
<u>LEDIPASVIR; SOFOSBUVIR - HARVONI</u>						
N 205834 001	10039779	Jan 30, 2034	DS DP	U-2369	D-177	Nov 15, 2022
	10039779	Jan 30, 2034	DS DP	U-2370	NPP	Apr 07, 2020
	10039779*PED	Jul 30, 2034			ODE-136	Apr 07, 2024
	10456414	Sep 14, 2032		DP		
	7964580	Mar 26, 2029	DS DP	U-1470	PED	Oct 07, 2024
	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				

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<u>LEDIPASVIR; SOFOSBUVIR - HARVONI</u>						
N 205834 001	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		D-177	Nov 15, 2022
	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				
	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		D-177	Nov 15, 2022
	7964580	Mar 26, 2029	DS DP U-1470		ODE-262	Aug 28, 2026
	7964580*PED	Sep 26, 2029			ODE-263	Aug 28, 2026
	8088368	May 12, 2030	DS DP		ODE-264	Aug 28, 2026
	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			

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<u>LEDIPASVIR; SOFOSBUVIR - HARVONI</u>						
N 212477 001	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	D-177	Nov 15, 2022
	7964580	Mar 26, 2029	DS DP U-1470		ODE-262	Aug 28, 2026
	7964580*PED	Sep 26, 2029			ODE-263	Aug 28, 2026
	8088368	May 12, 2030	DS DP		ODE-264	Aug 28, 2026
	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>LEFAMULIN ACETATE - XENLETA</u>						
N 211672 001	6753445	Jul 09, 2021	DS DP U-2619		NCE	Aug 19, 2024
	8071643	Jan 16, 2029	DS DP		GAIN	Aug 19, 2029
	8153689	Mar 19, 2028	DS DP			
	9120727	May 23, 2031	DS DP			
<u>LEFAMULIN ACETATE - XENLETA</u>						
N 211673 001	6753445	Jul 09, 2021	DS DP U-2619		NCE	Aug 19, 2024
	8071643	Jan 16, 2029	DS DP		GAIN	Aug 19, 2029
	8153689	Mar 19, 2028	DS DP			
<u>LENALIDOMIDE - REVLIMID</u>						
N 021880 001	6315720	Oct 23, 2020			I-796	May 28, 2022
	6561977	Oct 23, 2020			I-797	May 28, 2022
	6755784	Oct 23, 2020			ODE-131	Feb 22, 2024
	7189740	Apr 11, 2023			ODE-241	May 28, 2026
	7465800	Apr 27, 2027	DS DP		ODE-245	May 28, 2026
	7468363	Oct 07, 2023			ODE-49	Jun 05, 2020
	7468363	Oct 07, 2023			ODE-88	Feb 17, 2022
	7468363	Oct 07, 2023				
	7855217	Nov 24, 2024	DS DP			
	7968569	Oct 07, 2023			U-1984	
	8315886	Oct 23, 2020			U-1249	
	8404717	Apr 11, 2023			U-1982	
	8492406	Oct 07, 2023			U-2550	
	8530498	May 15, 2023			U-1984	
	8626531	Oct 23, 2020			U-1210	

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<u>LENALIDOMIDE - REVLIMID</u>						
N 021880 001	8648095	May 15, 2023		U-1984		
	8741929	Mar 08, 2028		U-1983		
	9056120	Apr 11, 2023		U-1982		
	9101621	May 15, 2023		U-1985		
	9101622	May 15, 2023		U-1986		
	9155730	May 15, 2023		U-2550		
	9393238	May 15, 2023		U-2550		
<u>LENALIDOMIDE - REVLIMID</u>						
N 021880 002	6315720	Oct 23, 2020		U-1210	I-796	May 28, 2022
	6561977	Oct 23, 2020		U-1210	I-797	May 28, 2022
	6755784	Oct 23, 2020		U-1210	ODE-131	Feb 22, 2024
	7189740	Apr 11, 2023		U-1982	ODE-241	May 28, 2026
	7465800	Apr 27, 2027	DS DP		ODE-245	May 28, 2026
	7468363	Oct 07, 2023		U-1983	ODE-49	Jun 05, 2020
	7468363	Oct 07, 2023		U-2550	ODE-88	Feb 17, 2022
	7468363	Oct 07, 2023		U-2551		
	7855217	Nov 24, 2024	DS DP			
	7968569	Oct 07, 2023		U-1984		
	8315886	Oct 23, 2020		U-1249		
	8404717	Apr 11, 2023		U-1982		
	8492406	Oct 07, 2023		U-2550		
	8530498	May 15, 2023		U-1984		
	8626531	Oct 23, 2020		U-1210		
	8648095	May 15, 2023		U-1984		
	8741929	Mar 08, 2028		U-1983		
	9056120	Apr 11, 2023		U-1982		
	9101621	May 15, 2023		U-1985		
	9101622	May 15, 2023		U-1986		
	9155730	May 15, 2023		U-2550		
	9393238	May 15, 2023		U-2550		
<u>LENALIDOMIDE - REVLIMID</u>						
N 021880 003	6315720	Oct 23, 2020		U-1210	I-796	May 28, 2022
	6561977	Oct 23, 2020		U-1210	I-797	May 28, 2022
	6755784	Oct 23, 2020		U-1210	ODE-131	Feb 22, 2024
	7189740	Apr 11, 2023		U-1982	ODE-241	May 28, 2026
	7465800	Apr 27, 2027	DS DP		ODE-245	May 28, 2026
	7468363	Oct 07, 2023		U-1983	ODE-49	Jun 05, 2020
	7468363	Oct 07, 2023		U-2550	ODE-88	Feb 17, 2022
	7468363	Oct 07, 2023		U-2551		
	7855217	Nov 24, 2024	DS DP			
	7968569	Oct 07, 2023		U-1984		
	8315886	Oct 23, 2020		U-1249		
	8404717	Apr 11, 2023		U-1982		
	8492406	Oct 07, 2023		U-2550		
	8530498	May 15, 2023		U-1984		
	8626531	Oct 23, 2020		U-1210		
	8648095	May 15, 2023		U-1984		
	8741929	Mar 08, 2028		U-1983		
	9056120	Apr 11, 2023		U-1982		
	9101621	May 15, 2023		U-1985		
	9101622	May 15, 2023		U-1986		
	9155730	May 15, 2023		U-2550		
	9393238	May 15, 2023		U-2550		
<u>LENALIDOMIDE - REVLIMID</u>						
N 021880 004	6315720	Oct 23, 2020		U-1210	I-796	May 28, 2022
	6561977	Oct 23, 2020		U-1210	I-797	May 28, 2022
	6755784	Oct 23, 2020		U-1210	ODE-131	Feb 22, 2024
	7189740	Apr 11, 2023		U-1982	ODE-241	May 28, 2026
	7465800	Apr 27, 2027	DS DP		ODE-245	May 28, 2026
	7468363	Oct 07, 2023		U-1983	ODE-49	Jun 05, 2020
	7468363	Oct 07, 2023		U-2550	ODE-88	Feb 17, 2022
	7468363	Oct 07, 2023		U-2551		
	7855217	Nov 24, 2024	DS DP			
	7968569	Oct 07, 2023		U-1984		
	8315886	Oct 23, 2020		U-1249		
	8404717	Apr 11, 2023		U-1982		
	8492406	Oct 07, 2023		U-2550		

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<u>LENALIDOMIDE - REVLIMID</u>						
N 021880 004	8530498	May 15, 2023	U-1984			
	8626531	Oct 23, 2020	U-1210			
	8648095	May 15, 2023	U-1984			
	8741929	Mar 08, 2028	U-1983			
	9056120	Apr 11, 2023	U-1982			
	9101621	May 15, 2023	U-1985			
	9101622	May 15, 2023	U-1986			
	9155730	May 15, 2023	U-2550			
	9393238	May 15, 2023	U-2550			
<u>LENALIDOMIDE - REVLIMID</u>						
N 021880 005	6315720	Oct 23, 2020	U-1210		I-796	May 28, 2022
	6561977	Oct 23, 2020	U-1210		I-797	May 28, 2022
	6755784	Oct 23, 2020	U-1210		ODE-131	Feb 22, 2024
	7189740	Apr 11, 2023	U-1982		ODE-241	May 28, 2026
	7465800	Apr 27, 2027	DS DP		ODE-245	May 28, 2026
	7468363	Oct 07, 2023	U-1983		ODE-49	Jun 05, 2020
	7468363	Oct 07, 2023	U-2550		ODE-88	Feb 17, 2022
	7468363	Oct 07, 2023	U-2551			
	7855217	Nov 24, 2024	DS DP			
	7968569	Oct 07, 2023	U-1984			
	8315886	Oct 23, 2020	U-1249			
	8404717	Apr 11, 2023	U-1982			
	8492406	Oct 07, 2023	U-2550			
	8530498	May 15, 2023	U-1984			
	8626531	Oct 23, 2020	U-1210			
	8648095	May 15, 2023	U-1984			
	8741929	Mar 08, 2028	U-1983			
	9056120	Apr 11, 2023	U-1982			
	9101621	May 15, 2023	U-1985			
	9101622	May 15, 2023	U-1986			
	9155730	May 15, 2023	U-2550			
	9393238	May 15, 2023	U-2550			
<u>LENALIDOMIDE - REVLIMID</u>						
N 021880 006	6315720	Oct 23, 2020	U-1210		I-796	May 28, 2022
	6561977	Oct 23, 2020	U-1210		I-797	May 28, 2022
	6755784	Oct 23, 2020	U-1210		ODE-131	Feb 22, 2024
	7189740	Apr 11, 2023	U-1982		ODE-241	May 28, 2026
	7465800	Apr 27, 2027	DS DP		ODE-245	May 28, 2026
	7468363	Oct 07, 2023	U-1983		ODE-49	Jun 05, 2020
	7468363	Oct 07, 2023	U-2550		ODE-88	Feb 17, 2022
	7468363	Oct 07, 2023	U-2551			
	7855217	Nov 24, 2024	DS DP			
	7968569	Oct 07, 2023	U-1984			
	8315886	Oct 23, 2020	U-1249			
	8404717	Apr 11, 2023	U-1982			
	8492406	Oct 07, 2023	U-2550			
	8530498	May 15, 2023	U-1984			
	8626531	Oct 23, 2020	U-1210			
	8648095	May 15, 2023	U-1984			
	8741929	Mar 08, 2028	U-1983			
	9056120	Apr 11, 2023	U-1982			
	9101621	May 15, 2023	U-1985			
	9101622	May 15, 2023	U-1986			
	9155730	May 15, 2023	U-2550			
	9393238	May 15, 2023	U-2550			
<u>LENVATINIB MESYLATE - LENVIMA</u>						
N 206947 001	10259791	Aug 26, 2035	DS		I-787	Aug 15, 2021
	10407393	Aug 26, 2035	DS		I-807	Sep 17, 2022
	7253286	Oct 19, 2021	DS DP		NCE	Feb 13, 2020
	7612208	Sep 19, 2026	DS DP		ODE-196	Aug 15, 2025
	9006256	Jul 27, 2027	U-1695		ODE-87	Feb 13, 2022
<u>LENVATINIB MESYLATE - LENVIMA</u>						
N 206947 002	10259791	Aug 26, 2035	DS		I-787	Aug 15, 2021
	10407393	Aug 26, 2035	DS		I-807	Sep 17, 2022
	7253286	Oct 19, 2021	DS DP		NCE	Feb 13, 2020
	7612208	Sep 19, 2026	DS DP		ODE-196	Aug 15, 2025

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<u>LENVATINIB MESYLATE - LENVIMA</u>						
N 206947	002 9006256	Jul 27, 2027	U-1695		ODE-87	Feb 13, 2022
<u>LESINURAD - ZURAMPIC</u>						
N 207988	001 10183012	Nov 26, 2028	U-2311		NCE	Dec 22, 2020
	8003681	Aug 25, 2025	DS			
	8084483	Aug 17, 2029	U-1801			
	8283369	Nov 26, 2028	U-1802			
	8283369	Nov 26, 2028	U-1804			
	8357713	Nov 26, 2028	DP U-1801			
	8357713	Nov 26, 2028	DP U-1802			
	8357713	Nov 26, 2028	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 7196086	May 22, 2024	DS DP		NCE	Nov 08, 2022
	8513255	May 22, 2024	DS DP		ODE-165	Nov 08, 2024
<u>LETERMOVIR - PREVYMIS</u>						
N 209939	002 7196086	May 22, 2024	DS DP		NCE	Nov 08, 2022
	8513255	May 22, 2024	DS DP		ODE-165	Nov 08, 2024
<u>LETERMOVIR - PREVYMIS</u>						
N 209940	001 7196086	May 22, 2024	DS DP		NCE	Nov 08, 2022
	8513255	May 22, 2024	DS DP		ODE-165	Nov 08, 2024
<u>LETERMOVIR - PREVYMIS</u>						
N 209940	002 7196086	May 22, 2024	DS DP		NCE	Nov 08, 2022
	8513255	May 22, 2024	DS DP		ODE-165	Nov 08, 2024
<u>LETROZOLE; RIBOCICLIB SUCCINATE - KISQALI FEMARA CO-PACK (COPACKAGED)</u>						
N 209935	001 8324225	Jun 17, 2028	DS DP		NCE	Mar 13, 2022
	8415355	Feb 19, 2031	DS DP			
	8685980	May 25, 2030	DS DP			
	8962630	Dec 09, 2029	U-2505			
	9193732	Nov 09, 2031	DS DP			
	9416136	Aug 20, 2029	U-2505			
	9868739	Nov 09, 2031	U-2505			
<u>LEUPROLIDE ACETATE - LUPRON DEPOT</u>						
N 020517	003 8815801	Jun 28, 2022	DP			
	8921326	Feb 05, 2031	DP U-1666			
<u>LEUPROLIDE ACETATE - ELIGARD</u>						
N 021343	001 6626870	Mar 27, 2020	DP			
<u>LEUPROLIDE ACETATE - ELIGARD</u>						
N 021379	001 6626870	Mar 27, 2020	DP			
	8470359	Oct 15, 2023	DS DP U-621			
	8840916	Nov 13, 2020	DP			
	9539333	Nov 13, 2020	DS DP U-621			
<u>LEUPROLIDE ACETATE - ELIGARD</u>						
N 021488	001 6626870	Mar 27, 2020	DP			
	8470359	Oct 15, 2023	DS DP U-621			
	8840916	Nov 13, 2020	DP			
	9539333	Nov 13, 2020	DS DP U-621			
<u>LEUPROLIDE ACETATE - ELIGARD</u>						
N 021731	001 6626870	Mar 27, 2020	DP			
	8470359	Oct 15, 2023	DS DP U-621			
	8840916	Nov 13, 2020	DP			
	9539333	Nov 13, 2020	DS DP U-621			
	9914802	Nov 13, 2020	DS DP U-1666			
<u>LEUPROLIDE ACETATE - LUTRATE DEPOT KIT</u>						
N 205054	001 9789064	Dec 15, 2020	DP			

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<u>LEVALBUTEROL HYDROCHLORIDE - XOPENEX</u>						
N 020837	001	6451289				Mar 21, 2021
<u>LEVALBUTEROL HYDROCHLORIDE - XOPENEX</u>						
N 020837	002	6451289				Mar 21, 2021
<u>LEVALBUTEROL HYDROCHLORIDE - XOPENEX</u>						
N 020837	003	6451289				Mar 21, 2021
<u>LEVALBUTEROL HYDROCHLORIDE - XOPENEX</u>						
N 020837	004	6451289		DP		Mar 21, 2021
<u>LEVALBUTEROL TARTRATE - XOPENEX HFA</u>						
N 021730	001	7256310		DS DP	U-636	Oct 08, 2024
		8765153		DP		Dec 08, 2023
<u>LEVETIRACETAM - KEPBRA</u>						
N 021035	001	8802142		DP		Jun 07, 2031
		8802142*PED				Dec 07, 2031
<u>LEVETIRACETAM - KEPBRA</u>						
N 021035	002	8802142		DP		Jun 07, 2031
		8802142*PED				Dec 07, 2031
<u>LEVETIRACETAM - KEPBRA</u>						
N 021035	003	8802142		DP		Jun 07, 2031
		8802142*PED				Dec 07, 2031
<u>LEVETIRACETAM - KEPBRA</u>						
N 021035	004	8802142		DP		Jun 07, 2031
		8802142*PED				Dec 07, 2031
<u>LEVETIRACETAM - KEPBRA XR</u>						
N 022285	001	7858122		DP		Sep 17, 2028
<u>LEVETIRACETAM - KEPBRA XR</u>						
N 022285	002	7858122		DP		Sep 17, 2028
<u>LEVETIRACETAM - ELEPSIA XR</u>						
N 204417	001	8163306		DP		Sep 03, 2027
		8425938		DP		Feb 22, 2026
		8431156		DP		Oct 31, 2027
		8470367		DP		Oct 31, 2027
		8535717		DP		Feb 22, 2026
<u>LEVETIRACETAM - ELEPSIA XR</u>						
N 204417	002	8163306		DP		Sep 03, 2027
		8425938		DP		Feb 22, 2026
		8431156		DP		Oct 31, 2027
		8470367		DP		Oct 31, 2027
		8535717		DP		Feb 22, 2026
<u>LEVETIRACETAM - SPRITAM</u>						
N 207958	001	9339489		DP	U-1850	Mar 14, 2034
		9669009			U-1850	Mar 14, 2034
		9669009			U-2021	Mar 14, 2034
		9669009			U-2022	Mar 14, 2034
<u>LEVETIRACETAM - SPRITAM</u>						
N 207958	002	9339489		DP	U-1850	Mar 14, 2034
		9669009			U-1850	Mar 14, 2034
		9669009			U-2021	Mar 14, 2034
		9669009			U-2022	Mar 14, 2034
<u>LEVETIRACETAM - SPRITAM</u>						
N 207958	003	9339489		DP	U-1850	Mar 14, 2034
		9669009			U-1850	Mar 14, 2034
		9669009			U-2021	Mar 14, 2034
		9669009			U-2022	Mar 14, 2034

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<u>LEVETIRACETAM - SPRITAM</u>						
N 207958	004	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>LEVOCARNITINE - LEVOCARNITINE</u>						
A 211676	001				CGT	Apr 13, 2020
<u>LEVOCARNITINE - LEVOCARNITINE SF</u>						
A 211676	002				CGT	Apr 13, 2020
<u>LEVOCARNITINE - CARNITOR</u>						
N 020182	001	6335369	Jan 18, 2021	U-433		
		6429230	Jan 18, 2021	U-433		
		6696493	Jan 18, 2021	U-433		
<u>LEVOCETIRIZINE DIHYDROCHLORIDE - XYZAL ALLERGY 24HR</u>						
N 209090	001	8633194	Oct 16, 2027	DP		
<u>LEVODOPA - INBRIJA</u>						
N 209184	001	6514482	Sep 19, 2020	DP U-2484	NP	Dec 21, 2021
		6613308	Sep 19, 2020	U-2484		
		6858199	Nov 04, 2021	U-2485		
		6921528	Jun 19, 2020	U-2485		
		6979437	Sep 19, 2020	U-2484		
		7146978	Apr 16, 2021	U-2486		
		7182961	Feb 22, 2024	DP		
		7384649	Nov 20, 2022	DP		
		7556798	Nov 04, 2021	U-2485		
		8404276	Mar 19, 2023	U-2484		
		8545878	Nov 16, 2032	DP		
		8586093	Mar 19, 2023	U-2484		
		8628754	Jun 19, 2020	U-2485		
		8685442	Nov 16, 2032	DP		
		8945612	Nov 16, 2032	DP		
		9155699	Mar 19, 2023	DP		
		9393210	Nov 16, 2032	DP		
		RE43711	Feb 03, 2029	U-2484		
<u>LEVOFLOXACIN - LEVAQUIN</u>						
N 021721	001	6806256	Feb 26, 2022	DP		
<u>LEVOLEUCOVORIN CALCIUM - FUSILEV</u>						
N 020140	001	6500829	Mar 07, 2022	DS DP		
<u>LEVOLEUCOVORIN CALCIUM - FUSILEV</u>						
N 020140	002	6500829	Mar 07, 2022	DS DP		
<u>LEVOLEUCOVORIN CALCIUM - FUSILEV</u>						
N 020140	003	6500829	Mar 07, 2022	DS DP		
<u>LEVOMILNACIPRAN HYDROCHLORIDE - FETZIMA</u>						
N 204168	001	8481598	Mar 02, 2031	U-839	M-249	Oct 07, 2022
		8865937	May 23, 2032	DS DP		
		RE43879	Jun 03, 2023	U-839		
<u>LEVOMILNACIPRAN HYDROCHLORIDE - FETZIMA</u>						
N 204168	002	8481598	Mar 02, 2031	U-839	M-249	Oct 07, 2022
		8865937	May 23, 2032	DS DP		
		RE43879	Jun 03, 2023	U-839		
<u>LEVOMILNACIPRAN HYDROCHLORIDE - FETZIMA</u>						
N 204168	003	8481598	Mar 02, 2031	U-839	M-249	Oct 07, 2022
		8865937	May 23, 2032	DS DP		
		RE43879	Jun 03, 2023	U-839		
<u>LEVOMILNACIPRAN HYDROCHLORIDE - FETZIMA</u>						
N 204168	004	8481598	Mar 02, 2031	U-839	M-249	Oct 07, 2022
		8865937	May 23, 2032	DS DP		
		RE43879	Jun 03, 2023	U-839		

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<u>LEVONORGESTREL - MIRENA</u>						
N 021225	001 9615965	Sep 16, 2029	DP U-2003			
	9668912	Apr 01, 2031	DP			
<u>LEVONORGESTREL - SKYLA</u>						
N 203159	001 7252839	Nov 13, 2023	DP			
	9615965	Sep 16, 2029	DP U-2003			
	9668912	Apr 01, 2031	DP			
<u>LEVONORGESTREL - LILETTA</u>						
N 206229	001 10028858	Mar 22, 2034	DP U-2348			
<u>LEVONORGESTREL - KYLEENA</u>						
N 208224	001 7252839	Nov 13, 2023	DP			
	9615965	Sep 16, 2029	DP U-2003			
	9668912	Apr 01, 2031	DP			
<u>LEVOTHYROXINE SODIUM - LEVOXYL</u>						
N 021301	001 6555581	Feb 15, 2022				
	7067148	Feb 15, 2022	DP			
	7101569	Oct 02, 2023			U-759	
<u>LEVOTHYROXINE SODIUM - LEVOXYL</u>						
N 021301	002 6555581	Feb 15, 2022				
	7067148	Feb 15, 2022	DP			
	7101569	Oct 02, 2023			U-759	
<u>LEVOTHYROXINE SODIUM - LEVOXYL</u>						
N 021301	003 6555581	Feb 15, 2022				
	7067148	Feb 15, 2022	DP			
	7101569	Oct 02, 2023			U-759	
<u>LEVOTHYROXINE SODIUM - LEVOXYL</u>						
N 021301	004 6555581	Feb 15, 2022				
	7067148	Feb 15, 2022	DP			
	7101569	Oct 02, 2023			U-759	
<u>LEVOTHYROXINE SODIUM - LEVOXYL</u>						
N 021301	005 6555581	Feb 15, 2022				
	7067148	Feb 15, 2022	DP			
	7101569	Oct 02, 2023			U-759	
<u>LEVOTHYROXINE SODIUM - LEVOXYL</u>						
N 021301	006 6555581	Feb 15, 2022				
	7067148	Feb 15, 2022	DP			
	7101569	Oct 02, 2023			U-759	
<u>LEVOTHYROXINE SODIUM - LEVOXYL</u>						
N 021301	007 6555581	Feb 15, 2022				
	7067148	Feb 15, 2022	DP			
	7101569	Oct 02, 2023			U-759	
<u>LEVOTHYROXINE SODIUM - LEVOXYL</u>						
N 021301	008 6555581	Feb 15, 2022				
	7067148	Feb 15, 2022	DP			
	7101569	Oct 02, 2023			U-759	
<u>LEVOTHYROXINE SODIUM - LEVOXYL</u>						
N 021301	009 6555581	Feb 15, 2022				
	7067148	Feb 15, 2022	DP			
	7101569	Oct 02, 2023			U-759	
<u>LEVOTHYROXINE SODIUM - LEVOXYL</u>						
N 021301	010 6555581	Feb 15, 2022				
	7067148	Feb 15, 2022	DP			
	7101569	Oct 02, 2023			U-759	
<u>LEVOTHYROXINE SODIUM - LEVOXYL</u>						
N 021301	011 6555581	Feb 15, 2022				
	7067148	Feb 15, 2022	DP			
	7101569	Oct 02, 2023			U-759	

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<u>LEVOTHYROXINE SODIUM - LEVOXYL</u>						
N 021301	012 655581	Feb 15, 2022				
	7067148	Feb 15, 2022	DP			
	7101569	Oct 02, 2023		U-759		
<u>LEVOTHYROXINE SODIUM - LEVO-T</u>						
N 021342	001 6399101	Mar 30, 2020				
<u>LEVOTHYROXINE SODIUM - LEVO-T</u>						
N 021342	002 6399101	Mar 30, 2020				
<u>LEVOTHYROXINE SODIUM - LEVO-T</u>						
N 021342	003 6399101	Mar 30, 2020				
<u>LEVOTHYROXINE SODIUM - LEVO-T</u>						
N 021342	004 6399101	Mar 30, 2020				
<u>LEVOTHYROXINE SODIUM - LEVO-T</u>						
N 021342	005 6399101	Mar 30, 2020				
<u>LEVOTHYROXINE SODIUM - LEVO-T</u>						
N 021342	006 6399101	Mar 30, 2020				
<u>LEVOTHYROXINE SODIUM - LEVO-T</u>						
N 021342	007 6399101	Mar 30, 2020				
<u>LEVOTHYROXINE SODIUM - LEVO-T</u>						
N 021342	008 6399101	Mar 30, 2020				
<u>LEVOTHYROXINE SODIUM - LEVO-T</u>						
N 021342	009 6399101	Mar 30, 2020				
<u>LEVOTHYROXINE SODIUM - LEVO-T</u>						
N 021342	010 6399101	Mar 30, 2020				
<u>LEVOTHYROXINE SODIUM - LEVO-T</u>						
N 021342	011 6399101	Mar 30, 2020				
<u>LEVOTHYROXINE SODIUM - TIROSINT</u>						
N 021924	002 7691411	Mar 14, 2024		DP		
	7723390	Mar 14, 2024		DP		
<u>LEVOTHYROXINE SODIUM - TIROSINT</u>						
N 021924	003 7691411	Mar 14, 2024		DP		
	7723390	Mar 14, 2024		DP		
<u>LEVOTHYROXINE SODIUM - TIROSINT</u>						
N 021924	004 7691411	Mar 14, 2024		DP		
	7723390	Mar 14, 2024		DP		
<u>LEVOTHYROXINE SODIUM - TIROSINT</u>						
N 021924	005 7691411	Mar 14, 2024		DP		
	7723390	Mar 14, 2024		DP		
<u>LEVOTHYROXINE SODIUM - TIROSINT</u>						
N 021924	006 7691411	Mar 14, 2024		DP		
	7723390	Mar 14, 2024		DP		
<u>LEVOTHYROXINE SODIUM - TIROSINT</u>						
N 021924	007 7691411	Mar 14, 2024		DP		
	7723390	Mar 14, 2024		DP		
<u>LEVOTHYROXINE SODIUM - TIROSINT</u>						
N 021924	008 7691411	Mar 14, 2024		DP		
	7723390	Mar 14, 2024		DP		
<u>LEVOTHYROXINE SODIUM - TIROSINT</u>						
N 021924	009 7691411	Mar 14, 2024		DP		
	7723390	Mar 14, 2024		DP		

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<u>LEVOTHYROXINE SODIUM - TIROSINT</u>						
N 021924 010	7691411	Mar 14, 2024	DP			
	7723390	Mar 14, 2024	DP			
<u>LEVOTHYROXINE SODIUM - TIROSINT</u>						
N 021924 011	7691411	Mar 14, 2024	DP			
	7723390	Mar 14, 2024	DP			
<u>LEVOTHYROXINE SODIUM - TIROSINT</u>						
N 021924 012	7691411	Mar 14, 2024	DP			
	7723390	Mar 14, 2024	DP			
<u>LEVOTHYROXINE SODIUM - TIROSINT</u>						
N 021924 013	7691411	Mar 14, 2024	DP			
	7723390	Mar 14, 2024	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			
<u>LEVOTHYROXINE SODIUM - TIROSINT-SOL</u>						
N 206977 002	10537538	Feb 28, 2037	DP			
<u>LEVOTHYROXINE SODIUM - TIROSINT-SOL</u>						
N 206977 003	10537538	Feb 28, 2037	DP			
<u>LEVOTHYROXINE SODIUM - TIROSINT-SOL</u>						
N 206977 004	10537538	Feb 28, 2037	DP			
<u>LEVOTHYROXINE SODIUM - TIROSINT-SOL</u>						
N 206977 005	10537538	Feb 28, 2037	DP			
<u>LEVOTHYROXINE SODIUM - TIROSINT-SOL</u>						
N 206977 006	10537538	Feb 28, 2037	DP			
<u>LEVOTHYROXINE SODIUM - TIROSINT-SOL</u>						
N 206977 007	10537538	Feb 28, 2037	DP			
<u>LEVOTHYROXINE SODIUM - TIROSINT-SOL</u>						
N 206977 008	10537538	Feb 28, 2037	DP			
<u>LEVOTHYROXINE SODIUM - TIROSINT-SOL</u>						
N 206977 009	10537538	Feb 28, 2037	DP			
<u>LEVOTHYROXINE SODIUM - TIROSINT-SOL</u>						
N 206977 010	10537538	Feb 28, 2037	DP			
<u>LEVOTHYROXINE SODIUM - TIROSINT-SOL</u>						
N 206977 011	10537538	Feb 28, 2037	DP			
<u>LEVOTHYROXINE SODIUM - TIROSINT-SOL</u>						
N 206977 012	10537538	Feb 28, 2037	DP			
<u>LEVOTHYROXINE SODIUM - LEVOTHYROXINE SODIUM</u>						
N 210632 001	10398669	Dec 01, 2036	DP			
	9782376	Dec 01, 2036	DP			

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<u>LEVOTHYROXINE SODIUM - LEVOTHYROXINE SODIUM</u>						
N 210632	002 10398669	Dec 01, 2036	DP			
	9782376	Dec 01, 2036	DP			
<u>LEVOTHYROXINE SODIUM - LEVOTHYROXINE SODIUM</u>						
N 210632	003 10398669	Dec 01, 2036	DP			
	9782376	Dec 01, 2036	DP			
<u>LIDOCAINE - ZTLIDO</u>						
N 207962	001 9283174	May 10, 2031	DP		NP	Feb 28, 2021
	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 - SYNERA</u>						
N 021623	001 6465709	Jul 07, 2020	DP			
<u>LIDOCAINE; TETRACAINE - PLIAGLIS</u>						
N 021717	001 10350180	Jan 14, 2031	DP			
<u>LIFITEGRAST - XIIDRA</u>						
N 208073	001 10124000	Nov 05, 2024		U-1900	NCE	Jul 11, 2021
	7314938	Mar 10, 2025	DS DP			
	7745460	Nov 05, 2024	DS DP	U-1880		
	7790743	Nov 05, 2024		U-1880		
	7928122	Nov 05, 2024	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			
	9216174	Nov 05, 2024	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		
	7304036	Aug 30, 2026	DS DP	U-1516		
	7371727	Jan 28, 2024	DS			
	7704947	Jan 28, 2024	DS DP			
	7745409	Jan 28, 2024	DS DP			
	8080526	Jan 28, 2024	DS DP			
	8110553	Jan 28, 2024		U-1278		
	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		
	7371727	Jan 28, 2024	DS			
	7704947	Jan 28, 2024	DS DP			
	7745409	Jan 28, 2024	DS DP			
	8080526	Jan 28, 2024	DS DP			
	8110553	Jan 28, 2024		U-1278		
	8748573	Oct 30, 2031		U-1515		
	8748573	Oct 30, 2031		U-1516		
	8802628	Oct 30, 2031	DP			
	8933030	Feb 17, 2031	DP			

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<u>LINACLOTIDE - LINZESS</u>						
N 202811 002	9708371	Aug 16, 2033	DP U-1515			
<u>LINACLOTIDE - LINZESS</u>						
N 202811 003	7304036	Aug 30, 2026	DS DP U-1516		NS	Jan 25, 2020
	7371727	Jan 28, 2024	DS			
	7704947	Jan 28, 2024	DS DP			
	7745409	Jan 28, 2024	DS DP			
	8080526	Jan 28, 2024	DS DP			
	8110553	Jan 28, 2024		U-1516		
	8933030	Feb 17, 2031	DP U-1516			
	9708371	Aug 16, 2033	DP U-1516			
<u>LINAGLIPTIN - TRADJENTA</u>						
N 201280 001	10034877	Aug 05, 2029		U-2347		
	7407955	May 02, 2025	DS DP			
	8119648	Aug 12, 2023		U-1270		
	8119648	Aug 12, 2023		U-774		
	8178541	Aug 12, 2023		U-1244		
	8178541	Aug 12, 2023		U-1245		
	8178541	Aug 12, 2023		U-1270		
	8178541	Aug 12, 2023		U-775		
	8673927	May 04, 2027		U-1503		
	8846695	Jun 04, 2030		U-1503	Y	
	8853156	Mar 05, 2031		U-1642		
	8883805	Nov 26, 2025	DP			
	9173859	May 04, 2027	DP U-1503			
	9173859	May 04, 2027	DP U-1768			
	9486526	Aug 05, 2029		U-1915		
<u>LINAGLIPTIN; METFORMIN HYDROCHLORIDE - JENTADUETO</u>						
N 201281 001	10022379	Apr 02, 2029		U-2339		
	7407955	May 02, 2025	DS DP			
	8119648	Aug 12, 2023		U-802		
	8178541	Aug 12, 2023	DP U-775			
	8673927	May 04, 2027		U-1503		
	8846695	Jun 04, 2030		U-1503		
	8883805	Nov 26, 2025	DP			
	9155705	May 21, 2030	DP			
	9173859	May 04, 2027	DP U-1503			
	9415016	Apr 02, 2029	DP			
<u>LINAGLIPTIN; METFORMIN HYDROCHLORIDE - JENTADUETO</u>						
N 201281 002	10022379	Apr 02, 2029		U-2339		
	7407955	May 02, 2025	DS DP			
	8119648	Aug 12, 2023		U-802		
	8178541	Aug 12, 2023	DP U-775			
	8673927	May 04, 2027		U-1503		
	8846695	Jun 04, 2030		U-1503		
	8883805	Nov 26, 2025	DP			
	9155705	May 21, 2030	DP			
	9173859	May 04, 2027	DP U-1503			
	9415016	Apr 02, 2029	DP			
<u>LINAGLIPTIN; METFORMIN HYDROCHLORIDE - JENTADUETO</u>						
N 201281 003	10022379	Apr 02, 2029		U-2339		
	7407955	May 02, 2025	DS DP			
	8119648	Aug 12, 2023		U-802		
	8178541	Aug 12, 2023	DP U-775			
	8673927	May 04, 2027		U-1503		
	8846695	Jun 04, 2030		U-1503	Y	
	8883805	Nov 26, 2025	DP			
	9155705	May 21, 2030	DP			
	9173859	May 04, 2027	DP U-1503			
	9415016	Apr 02, 2029	DP			
<u>LINAGLIPTIN; METFORMIN HYDROCHLORIDE - JENTADUETO XR</u>						
N 208026 001	10022379	Apr 02, 2029		U-2339		
	6488962	Jun 20, 2020	DP			
	7407955	May 02, 2025	DS DP			
	8119648	Aug 12, 2023		U-802		

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<u>LINAGLIPTIN; METFORMIN HYDROCHLORIDE - JENTADUETO XR</u>						
N 208026 001	8178541	Aug 12, 2023	DP U-1853			
	8673927	May 04, 2027	U-1503			
	8883805	Nov 26, 2025	DP			
	9155705	May 21, 2030	DP			
	9173859	May 04, 2027	DP U-1503			
	9415016	Apr 02, 2029	DP			
	9555001	Mar 06, 2033	DP U-1967			
	9555001	Mar 06, 2033	DP U-1968			
<u>LINAGLIPTIN; METFORMIN HYDROCHLORIDE - JENTADUETO XR</u>						
N 208026 002	10022379	Apr 02, 2029	U-2339			
	6488962	Jun 20, 2020	DP			
	7407955	May 02, 2025	DS DP			
	8119648	Aug 12, 2023	U-802			
	8178541	Aug 12, 2023	DP U-1853			
	8673927	May 04, 2027	U-1503			
	8883805	Nov 26, 2025	DP			
	9155705	May 21, 2030	DP			
	9173859	May 04, 2027	DP U-1503			
	9415016	Apr 02, 2029	DP			
	9555001	Mar 06, 2033	DP U-1967			
	9555001	Mar 06, 2033	DP U-1968			
<u>LINEZOLID - ZYVOX</u>						
N 021130 001	6514529	Mar 15, 2021	DP			
	6559305	Jan 29, 2021	DS			
<u>LINEZOLID - ZYVOX</u>						
N 021130 002	6514529	Mar 15, 2021	DP			
	6559305	Jan 29, 2021	DS			
<u>LINEZOLID - ZYVOX</u>						
N 021131 001	6559305	Jan 29, 2021	DS			
<u>LINEZOLID - ZYVOX</u>						
N 021131 002	6559305	Jan 29, 2021	DS			
	6559305*PED	Jul 29, 2021				
<u>LINEZOLID - ZYVOX</u>						
N 021131 003	6559305	Jan 29, 2021	DS			
	6559305*PED	Jul 29, 2021				
<u>LINEZOLID - ZYVOX</u>						
N 021132 001	6559305	Jan 29, 2021	DS			
<u>LIRAGLUTIDE RECOMBINANT - VICTOZA</u>						
N 022341 001	6268343	Aug 22, 2022	DS DP U-968		I-750	Aug 25, 2020
	6268343*PED	Feb 22, 2023			NPP	Jun 17, 2022
	7762994	May 23, 2024	DP		PED	Dec 17, 2022
	7762994*PED	Nov 23, 2024				
	8114833	Aug 13, 2025	DS DP			
	8114833*PED	Feb 13, 2026				
	8579869	Jun 30, 2023	DP			
	8579869*PED	Dec 30, 2023				
	8846618	Jun 27, 2022	DP			
	8846618*PED	Dec 27, 2022				
	9265893	Sep 23, 2032	DP			
	9265893*PED	Mar 23, 2033				
	9968659	Jan 09, 2037	U-2313			
	9968659*PED	Jul 09, 2037				
	RE41956	Jan 21, 2021	DP			
	RE41956*PED	Jul 21, 2021				
<u>LIRAGLUTIDE RECOMBINANT - 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			
	6268343	Aug 22, 2022	DS DP U-1255			
	6268343*PED	Feb 22, 2023				

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<u>LIRAGLUTIDE RECOMBINANT - SAXENDA</u>						
N 206321 001	6899699	Jan 01, 2022	DP			
	6899699*PED	Jul 01, 2022				
	7762994	May 23, 2024	DP			
	7762994*PED	Nov 23, 2024				
	8114833	Aug 13, 2025	DP			
	8114833*PED	Feb 13, 2026				
	8579869	Jun 30, 2023	DP			
	8579869*PED	Dec 30, 2023				
	8672898	Jan 02, 2022	DP			
	8672898*PED	Jul 02, 2022				
	8684969	Oct 20, 2025	DP			
	8684969*PED	Apr 20, 2026				
	8846618	Jun 27, 2022	DP			
	8846618*PED	Dec 27, 2022				
	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			
	9457154*PED	Mar 27, 2028				
	9486588	Jan 02, 2022	DP			
	9486588*PED	Jul 02, 2022				
	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>LISDEXAMFETAMINE DIMESYLATE - VYVANSE</u>						
N 021977 001	7105486	Feb 24, 2023		U-727		
	7223735	Feb 24, 2023	DP			
	7655630	Feb 24, 2023	DS			
	7659253	Feb 24, 2023	DS DP	U-727		
	7659254	Feb 24, 2023			U-1034	
	7662787	Feb 24, 2023	DS			
	7662788	Feb 24, 2023			U-727	
	7671030	Feb 24, 2023	DP	U-727		
	7671031	Feb 24, 2023			U-727	
	7674774	Feb 24, 2023	DP	U-842		
	7678770	Feb 24, 2023			U-842	
	7678771	Feb 24, 2023	DP	U-842		
	7687466	Feb 24, 2023	DP			
	7687467	Feb 24, 2023	DP	U-842		
	7700561	Feb 24, 2023	DP			
	7713936	Feb 24, 2023			U-727	
	7718619	Feb 24, 2023	DP	U-842		
	7723305	Feb 24, 2023	DP	U-842		
<u>LISDEXAMFETAMINE DIMESYLATE - VYVANSE</u>						
N 021977 002	7105486	Feb 24, 2023		U-727		
	7223735	Feb 24, 2023	DP			
	7655630	Feb 24, 2023	DS			
	7659253	Feb 24, 2023	DS DP	U-727		
	7659254	Feb 24, 2023			U-1034	
	7662787	Feb 24, 2023	DS			
	7662788	Feb 24, 2023			U-727	
	7671030	Feb 24, 2023	DP	U-727		
	7671031	Feb 24, 2023			U-727	
	7674774	Feb 24, 2023	DP	U-842		
	7678770	Feb 24, 2023			U-842	

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<u>LISDEXAMFETAMINE DIMESYLATE - VYVANSE</u>						
N 021977 002	7678771	Feb 24, 2023	DP U-842			
	7687466	Feb 24, 2023	DP			
	7687467	Feb 24, 2023	DP U-842			
	7700561	Feb 24, 2023	DP			
	7713936	Feb 24, 2023	U-727			
	7718619	Feb 24, 2023	DP U-842			
	7723305	Feb 24, 2023	DP U-842			
<u>LISDEXAMFETAMINE DIMESYLATE - VYVANSE</u>						
N 021977 003	7105486	Feb 24, 2023	U-727			
	7223735	Feb 24, 2023	DP			
	7655630	Feb 24, 2023	DS			
	7659253	Feb 24, 2023	DS DP U-727			
	7659254	Feb 24, 2023	U-1034			
	7662787	Feb 24, 2023	DS			
	7662788	Feb 24, 2023	U-727			
	7671030	Feb 24, 2023	DP U-727			
	7671031	Feb 24, 2023	U-727			
	7674774	Feb 24, 2023	DP U-842			
	7678770	Feb 24, 2023	U-842			
	7678771	Feb 24, 2023	DP U-842			
	7687466	Feb 24, 2023	DP			
	7687467	Feb 24, 2023	DP U-842			
	7700561	Feb 24, 2023	DP			
	7713936	Feb 24, 2023	U-727			
	7718619	Feb 24, 2023	DP U-842			
	7723305	Feb 24, 2023	DP U-842			
<u>LISDEXAMFETAMINE DIMESYLATE - VYVANSE</u>						
N 021977 004	7105486	Feb 24, 2023	U-727			
	7105486	Feb 24, 2023	U-842			
	7223735	Feb 24, 2023	DP			
	7655630	Feb 24, 2023	DS			
	7659253	Feb 24, 2023	DS DP U-727			
	7659254	Feb 24, 2023	U-1034			
	7662787	Feb 24, 2023	DS			
	7662788	Feb 24, 2023	U-727			
	7671030	Feb 24, 2023	DP U-727			
	7671031	Feb 24, 2023	U-727			
	7674774	Feb 24, 2023	DP U-842			
	7678770	Feb 24, 2023	U-842			
	7678771	Feb 24, 2023	DP U-842			
	7687466	Feb 24, 2023	DP			
	7687467	Feb 24, 2023	DP U-842			
	7700561	Feb 24, 2023	DP			
	7713936	Feb 24, 2023	U-727			
	7718619	Feb 24, 2023	DP U-842			
	7723305	Feb 24, 2023	DP U-842			
<u>LISDEXAMFETAMINE DIMESYLATE - VYVANSE</u>						
N 021977 005	7105486	Feb 24, 2023	U-842			
	7223735	Feb 24, 2023	DP			
	7655630	Feb 24, 2023	DS			
	7659253	Feb 24, 2023	DS DP U-727			
	7659254	Feb 24, 2023	U-1034			
	7662787	Feb 24, 2023	DS			
	7662788	Feb 24, 2023	U-727			
	7671030	Feb 24, 2023	DP U-727			
	7671031	Feb 24, 2023	U-727			
	7674774	Feb 24, 2023	DP U-842			
	7678770	Feb 24, 2023	U-842			
	7678771	Feb 24, 2023	DP U-842			
	7687466	Feb 24, 2023	DP			
	7687467	Feb 24, 2023	DP U-842			
	7700561	Feb 24, 2023	DP			
	7713936	Feb 24, 2023	U-727			
	7718619	Feb 24, 2023	DP U-842			
	7723305	Feb 24, 2023	DP U-842			

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<u>LISDEXAMFETAMINE DIMESYLATE - VYVANSE</u>						
N 021977 006	7105486	Feb 24, 2023	U-727			
	7105486	Feb 24, 2023	U-842			
	7223735	Feb 24, 2023	DP			
	7655630	Feb 24, 2023	DS			
	7659253	Feb 24, 2023	DS DP U-727			
	7659254	Feb 24, 2023	U-1034			
	7662787	Feb 24, 2023	DS			
	7662788	Feb 24, 2023	U-727			
	7671030	Feb 24, 2023	DP U-727			
	7671031	Feb 24, 2023	U-727			
	7674774	Feb 24, 2023	DP U-842			
	7678770	Feb 24, 2023	U-842			
	7678771	Feb 24, 2023	DP U-842			
	7687466	Feb 24, 2023	DP			
	7687467	Feb 24, 2023	DP U-842			
	7700561	Feb 24, 2023	DP			
	7713936	Feb 24, 2023	U-727			
	7718619	Feb 24, 2023	DP U-842			
	7723305	Feb 24, 2023	DP U-842			
<u>LISDEXAMFETAMINE DIMESYLATE - VYVANSE</u>						
N 021977 007	7223735	Feb 24, 2023	DP			
	7655630	Feb 24, 2023	DS			
	7659253	Feb 24, 2023	DS DP U-727			
	7659254	Feb 24, 2023	U-1034			
	7662787	Feb 24, 2023	DS			
	7662788	Feb 24, 2023	U-727			
	7671030	Feb 24, 2023	DP U-727			
	7671031	Feb 24, 2023	U-727			
	7674774	Feb 24, 2023	DP U-842			
	7678770	Feb 24, 2023	U-842			
	7678771	Feb 24, 2023	DP			
	7687466	Feb 24, 2023	DP			
	7687467	Feb 24, 2023	DP U-842			
	7700561	Feb 24, 2023	DP			
	7713936	Feb 24, 2023	U-727			
	7718619	Feb 24, 2023	DP U-842			
<u>LISDEXAMFETAMINE DIMESYLATE - VYVANSE</u>						
N 208510 001	7105486	Feb 24, 2023	U-727			
	7223735	Feb 24, 2023	DP			
	7655630	Feb 24, 2023	DS DP			
	7659253	Feb 24, 2023	DS DP U-727			
	7659254	Feb 24, 2023	U-727			
	7662787	Feb 24, 2023	DS			
	7662788	Feb 24, 2023	U-727			
	7671030	Feb 24, 2023	DP U-727			
	7671031	Feb 24, 2023	U-727			
	7674774	Feb 24, 2023	DP U-727			
	7678770	Feb 24, 2023	U-727			
	7678771	Feb 24, 2023	DP U-727			
	7687466	Feb 24, 2023	DP			
	7687467	Feb 24, 2023	DP U-727			
	7713936	Feb 24, 2023	U-727			
	7718619	Feb 24, 2023	DP U-727			
	7723305	Feb 24, 2023	DP U-727			
<u>LISDEXAMFETAMINE DIMESYLATE - VYVANSE</u>						
N 208510 002	7105486	Feb 24, 2023	U-727			
	7223735	Feb 24, 2023	DP			
	7655630	Feb 24, 2023	DS DP			
	7659253	Feb 24, 2023	DS DP U-727			
	7659254	Feb 24, 2023	U-727			
	7662787	Feb 24, 2023	DS			
	7662788	Feb 24, 2023	U-727			
	7671030	Feb 24, 2023	DP U-727			
	7671031	Feb 24, 2023	U-727			
	7674774	Feb 24, 2023	DP U-727			
	7678770	Feb 24, 2023	U-727			
	7678771	Feb 24, 2023	DP U-727			

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<u>LISDEXAMFETAMINE DIMESYLATE - VYVANSE</u>						
N 208510 002	7687466	Feb 24, 2023	DP			
	7687467	Feb 24, 2023	DP U-727			
	7713936	Feb 24, 2023	U-727			
	7718619	Feb 24, 2023	DP U-727			
	7723305	Feb 24, 2023	DP U-727			
<u>LISDEXAMFETAMINE DIMESYLATE - VYVANSE</u>						
N 208510 003	7105486	Feb 24, 2023			U-727	
	7223735	Feb 24, 2023	DP			
	7655630	Feb 24, 2023	DS DP			
	7659253	Feb 24, 2023	DS DP U-727			
	7659254	Feb 24, 2023	U-727			
	7662787	Feb 24, 2023	DS			
	7662788	Feb 24, 2023	U-727			
	7671030	Feb 24, 2023	DP U-727			
	7671031	Feb 24, 2023	U-727			
	7674774	Feb 24, 2023	DP U-727			
	7678770	Feb 24, 2023	U-727			
	7678771	Feb 24, 2023	DP U-727			
	7687466	Feb 24, 2023	DP			
	7687467	Feb 24, 2023	DP U-727			
	7713936	Feb 24, 2023	U-727			
	7718619	Feb 24, 2023	DP U-727			
	7723305	Feb 24, 2023	DP U-727			
<u>LISDEXAMFETAMINE DIMESYLATE - VYVANSE</u>						
N 208510 004	7105486	Feb 24, 2023			U-727	
	7223735	Feb 24, 2023	DP			
	7655630	Feb 24, 2023	DS DP			
	7659253	Feb 24, 2023	DS DP U-727			
	7659254	Feb 24, 2023	U-727			
	7662787	Feb 24, 2023	DS			
	7662788	Feb 24, 2023	U-727			
	7671030	Feb 24, 2023	DP U-727			
	7671031	Feb 24, 2023	U-727			
	7674774	Feb 24, 2023	DP U-727			
	7678770	Feb 24, 2023	U-727			
	7678771	Feb 24, 2023	DP U-727			
	7687466	Feb 24, 2023	DP			
	7687467	Feb 24, 2023	DP U-727			
	7713936	Feb 24, 2023	U-727			
	7718619	Feb 24, 2023	DP U-727			
	7723305	Feb 24, 2023	DP U-727			
<u>LISDEXAMFETAMINE DIMESYLATE - VYVANSE</u>						
N 208510 005	7105486	Feb 24, 2023			U-727	
	7223735	Feb 24, 2023	DP			
	7655630	Feb 24, 2023	DS DP			
	7659253	Feb 24, 2023	DS DP U-727			
	7659254	Feb 24, 2023	U-727			
	7662787	Feb 24, 2023	DS			
	7662788	Feb 24, 2023	U-727			
	7671030	Feb 24, 2023	DP U-727			
	7671031	Feb 24, 2023	U-727			
	7674774	Feb 24, 2023	DP U-727			
	7678770	Feb 24, 2023	U-727			
	7678771	Feb 24, 2023	DP U-727			
	7687466	Feb 24, 2023	DP			
	7687467	Feb 24, 2023	DP U-727			
	7713936	Feb 24, 2023	U-727			
	7718619	Feb 24, 2023	DP U-727			
	7723305	Feb 24, 2023	DP U-727			
<u>LISDEXAMFETAMINE DIMESYLATE - VYVANSE</u>						
N 208510 006	7105486	Feb 24, 2023			U-727	
	7223735	Feb 24, 2023	DP			
	7655630	Feb 24, 2023	DS DP			
	7659253	Feb 24, 2023	DS DP U-727			
	7659254	Feb 24, 2023	U-727			
	7662787	Feb 24, 2023	DS			

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<u>LISDEXAMFETAMINE DIMESYLATE - VYVANSE</u>						
N 208510	006	7662788	Feb 24, 2023			
		7671030	Feb 24, 2023			
		7671031	Feb 24, 2023			
		7674774	Feb 24, 2023	DP	U-727	
		7678770	Feb 24, 2023			
		7678771	Feb 24, 2023	DP	U-727	
		7687466	Feb 24, 2023	DP		
		7687467	Feb 24, 2023	DP	U-727	
		7713936	Feb 24, 2023			
		7718619	Feb 24, 2023	DP	U-727	
		7723305	Feb 24, 2023	DP	U-727	
<u>LISINOPRIL - OBRELIS</u>						
N 208401	001	10039800	Nov 06, 2035			
		10039800	Nov 06, 2035			
		10039800	Nov 06, 2035			
		10039800	Nov 06, 2035			
		10039800	Nov 06, 2035			
		10039800	Nov 06, 2035			
		10039800	Nov 06, 2035			
		10265370	Nov 06, 2035	DP		
		10406199	Nov 06, 2035			
		10406199	Nov 06, 2035			
		10406199	Nov 06, 2035			
		10406199	Nov 06, 2035			
		10406199	Nov 06, 2035			
		10406199	Nov 06, 2035			
		10406199	Nov 06, 2035			
		10406199	Nov 06, 2035			
		10406199	Nov 06, 2035			
		10406199	Nov 06, 2035			
		10406199	Nov 06, 2035			
		9463183	Nov 06, 2035	DP		
		9616096	Nov 06, 2035			
		9616096	Nov 06, 2035			
		9616096	Nov 06, 2035			
		9616096	Nov 06, 2035			
		9616096	Nov 06, 2035			
		9616096	Nov 06, 2035			
		9616096	Nov 06, 2035			
		9616096	Nov 06, 2035			
		9616096	Nov 06, 2035			
		9616096	Nov 06, 2035			
		9814751	Nov 06, 2035	DP		
<u>LIXISENATIDE - ADLYXIN</u>						
N 208471	001	10028910	Nov 11, 2030	DP	NCE	Jul 27, 2021
		10201663	Mar 10, 2034	DP		
		8475414	Dec 28, 2030	DP	U-1881	
		8882721	Jun 28, 2031	DP		
		8915888	Jun 08, 2030	DP	U-1881	
		9072836	Mar 15, 2032	DP		
		9084853	Oct 05, 2031	DP		
		9308329	Dec 28, 2030	DP	U-1881	
		9408893	Aug 27, 2032		U-1894	
		9440029	Jan 30, 2032	DP		
		9511193	Jan 19, 2032	DP		
		9707176	Nov 11, 2030	DP		
		9821032	May 09, 2032		U-2200	
		9855388	Apr 24, 2029	DP	U-1881	
		9981013	Aug 30, 2030		U-2297	
		RE45313	Jul 12, 2020	DS DP		
<u>LIXISENATIDE - ADLYXIN</u>						
N 208471	002	10028910	Nov 11, 2030	DP	NCE	Jul 27, 2021
		10201663	Mar 10, 2034	DP		
		8475414	Dec 28, 2030	DP	U-1881	
		8882721	Jun 28, 2031	DP		
		8915888	Jun 08, 2030	DP	U-1881	
		9072836	Mar 15, 2032	DP		
		9084853	Oct 05, 2031	DP		
		9308329	Dec 28, 2030	DP	U-1881	
		9408893	Aug 27, 2032		U-1894	
		9440029	Jan 30, 2032	DP		
		9511193	Jan 19, 2032	DP		
		9707176	Nov 11, 2030	DP		
		9821032	May 09, 2032		U-2200	

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<u>LIXISENATIDE - ADLYXIN</u>						
N 208471	002	9855388				
		Apr 24, 2029	DP	U-1881		
		9981013		U-2297		
		Aug 30, 2030				
		RE45313	DS	DP		
		Jul 12, 2020				
<u>LOFEXIDINE HYDROCHLORIDE - LUCEMYRA</u>						
N 209229	001				NCE	May 16, 2023
<u>LOMITAPIDE MESYLATE - JUXTAPID</u>						
N 203858	001	10016404				
		Mar 07, 2025		U-1316		
		5712279				
		Feb 21, 2020	DS	DP	U-1317	
		7932268			U-1316	
		Aug 19, 2027			U-1316	
		8618135			U-1316	
		Mar 07, 2025			U-1316	
		9265758			U-1316	
		Mar 07, 2025			U-1316	
		9364470			U-1851	
		Mar 07, 2025			U-1316	
		9433617			U-1316	
		Mar 07, 2025			U-1316	
		861622			U-1316	
		Mar 07, 2025			U-1316	
<u>LOMITAPIDE MESYLATE - JUXTAPID</u>						
N 203858	002	10016404				
		Mar 07, 2025		U-1316		
		5712279				
		Feb 21, 2020	DS	DP	U-1317	
		7932268			U-1316	
		Aug 19, 2027			U-1316	
		8618135			U-1316	
		Mar 07, 2025			U-1316	
		9265758			U-1316	
		Mar 07, 2025			U-1316	
		9364470			U-1851	
		Mar 07, 2025			U-1316	
		9433617			U-1316	
		Mar 07, 2025			U-1316	
		861622			U-1316	
		Mar 07, 2025			U-1316	
<u>LOMITAPIDE MESYLATE - JUXTAPID</u>						
N 203858	003	10016404				
		Mar 07, 2025		U-1316		
		5712279				
		Feb 21, 2020	DS	DP	U-1317	
		7932268			U-1316	
		Aug 19, 2027			U-1316	
		8618135			U-1316	
		Mar 07, 2025			U-1316	
		9265758			U-1316	
		Mar 07, 2025			U-1316	
		9364470			U-1851	
		Mar 07, 2025			U-1316	
		9433617			U-1316	
		Mar 07, 2025			U-1316	
		861622			U-1316	
		Mar 07, 2025			U-1316	
<u>LOMITAPIDE MESYLATE - JUXTAPID</u>						
N 203858	004	10016404				
		Mar 07, 2025		U-1316		
		5712279				
		Feb 21, 2020	DS	DP	U-1317	
		7932268			U-1316	
		Aug 19, 2027			U-1316	
		8618135			U-1316	
		Mar 07, 2025			U-1316	
		9265758			U-1316	
		Mar 07, 2025			U-1316	
		9364470			U-1851	
		Mar 07, 2025			U-1316	
		9433617			U-1316	
		Mar 07, 2025			U-1316	
		861622			U-1316	
		Mar 07, 2025			U-1316	
<u>LOMITAPIDE MESYLATE - JUXTAPID</u>						
N 203858	005	10016404				
		Mar 07, 2025		U-1316		
		5712279				
		Feb 21, 2020	DS	DP	U-1317	
		7932268			U-1316	
		Aug 19, 2027			U-1316	
		8618135			U-1316	
		Mar 07, 2025			U-1316	
		9265758			U-1316	
		Mar 07, 2025			U-1316	
		9364470			U-1851	
		Mar 07, 2025			U-1316	
		9433617			U-1316	
		Mar 07, 2025			U-1316	
		861622			U-1316	
		Mar 07, 2025			U-1316	
<u>LOMITAPIDE MESYLATE - JUXTAPID</u>						
N 203858	006	10016404				
		Mar 07, 2025		U-1316		
		5712279				
		Feb 21, 2020	DS	DP	U-1317	
		7932268			U-1316	
		Aug 19, 2027			U-1316	
		8618135			U-1316	
		Mar 07, 2025			U-1316	
		9265758			U-1316	
		Mar 07, 2025			U-1316	
		9364470			U-1851	
		Mar 07, 2025			U-1316	
		9433617			U-1316	
		Mar 07, 2025			U-1316	
		861622			U-1316	
		Mar 07, 2025			U-1316	
<u>LOPERAMIDE HYDROCHLORIDE - IMODIUM A-D EZ CHEWS</u>						
N 020448	001	6814978				
		Aug 26, 2021	DP			

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<u>LOPINAVIR; RITONAVIR - KALETRA</u>						
N 021226	001	7141593	May 22, 2020	DP		
		7432294	May 22, 2020	DP		
<u>LOPINAVIR; RITONAVIR - KALETRA</u>						
N 021251	001	6911214	Nov 28, 2021	DP	U-895	
		8501219	Nov 28, 2021	DP		
<u>LOPINAVIR; RITONAVIR - KALETRA</u>						
N 021906	001	7364752	Nov 10, 2020	DP	U-688	
		8025899	Dec 14, 2027	DP		
		8025899*PED	Jun 14, 2028			
		8268349	Aug 25, 2024	DP		
		8309613	Dec 24, 2024		U-688	
		8377952	Oct 22, 2027		U-1372	
		8377952*PED	Apr 22, 2028			
		8399015	Aug 25, 2024	DP		
		8399015*PED	Feb 25, 2025			
		8470347	Sep 17, 2026	DP		
		8470347*PED	Mar 17, 2027			
		8691878	Aug 25, 2024		U-1513	
		8691878*PED	Feb 25, 2025			
<u>LOPINAVIR; RITONAVIR - KALETRA</u>						
N 021906	002	7364752	Nov 10, 2020	DP	U-688	
		8025899	Dec 14, 2027	DP		
		8025899*PED	Jun 14, 2028			
		8268349	Aug 25, 2024	DP		
		8309613	Dec 24, 2024		U-688	
		8377952	Oct 22, 2027		U-1372	
		8377952*PED	Apr 22, 2028			
		8399015	Aug 25, 2024	DP		
		8399015*PED	Feb 25, 2025			
		8470347	Sep 17, 2026	DP		
		8470347*PED	Mar 17, 2027			
		8691878	Aug 25, 2024		U-1513	
		8691878*PED	Feb 25, 2025			
<u>LORCASERIN HYDROCHLORIDE - BELVIO</u>						
N 022529	001	6953787	Apr 10, 2023	DS DP	U-1252	
		6953787	Apr 10, 2023	DS DP	U-1253	
		6953787	Apr 10, 2023	DS DP	U-1254	
		6953787	Apr 10, 2023	DS DP	U-1255	
		7514422	Apr 10, 2023		U-1252	
		7514422	Apr 10, 2023		U-1253	
		7514422	Apr 10, 2023		U-1254	
		7514422	Apr 10, 2023		U-1255	
		7977329	Apr 10, 2023	DS DP	U-1252	
		7977329	Apr 10, 2023	DS DP	U-1253	
		7977329	Apr 10, 2023	DS DP	U-1254	
		7977329	Apr 10, 2023	DS DP	U-1255	
		8168624	Apr 18, 2029	DS DP		
		8207158	Apr 10, 2023		U-1252	
		8207158	Apr 10, 2023		U-1253	
		8207158	Apr 10, 2023		U-1254	
		8207158	Apr 10, 2023		U-1255	
		8273734	Apr 10, 2023		U-1254	
		8273734	Apr 10, 2023		U-1255	
		8367657	Apr 10, 2023	DS DP	U-1252	
		8367657	Apr 10, 2023	DS DP	U-1253	
		8367657	Apr 10, 2023	DS DP	U-1254	
		8367657	Apr 10, 2023	DS DP	U-1255	
		8546379	Apr 10, 2023	DS DP	U-1252	
		8546379	Apr 10, 2023	DS DP	U-1253	
		8546379	Apr 10, 2023	DS DP	U-1254	
		8546379	Apr 10, 2023	DS DP	U-1255	
		8575149	Apr 10, 2023		U-1452	
		8697686	Dec 20, 2025	DS DP		
		8946207	Jun 16, 2024	DP		
		8980881	Dec 20, 2025		U-1252	
		8980881	Dec 20, 2025		U-1253	

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<u>LORCASERIN HYDROCHLORIDE - BELVIO</u>						
N 022529	001	8980881	Dec 20, 2025		U-1254	
		8980881	Dec 20, 2025		U-1255	
		8999970	Feb 07, 2033		U-1688	
		8999970	Feb 07, 2033		U-1689	
		8999970	Feb 07, 2033		U-1692	
		9169213	Dec 06, 2032		U-1762	
		9169213	Dec 06, 2032		U-1763	
		9169213	Dec 06, 2032		U-1764	
		9169213	Dec 06, 2032		U-1765	
		9770455	Aug 31, 2031		U-2110	
<u>LORCASERIN HYDROCHLORIDE - BELVIO XR</u>						
N 208524	001	10226471	Aug 31, 2031	DP		
		10463676	Aug 31, 2031		U-2661	
		6953787	Apr 10, 2023	DS DP	U-1252	
		6953787	Apr 10, 2023	DS DP	U-1253	
		6953787	Apr 10, 2023	DS DP	U-1254	
		6953787	Apr 10, 2023	DS DP	U-1255	
		7514422	Apr 10, 2023		U-1252	
		7514422	Apr 10, 2023		U-1253	
		7514422	Apr 10, 2023		U-1254	
		7514422	Apr 10, 2023		U-1255	
		7977329	Apr 10, 2023	DS DP	U-1252	
		7977329	Apr 10, 2023	DS DP	U-1253	
		7977329	Apr 10, 2023	DS DP	U-1254	
		7977329	Apr 10, 2023	DS DP	U-1255	
		8168624	Apr 18, 2029	DS DP		
		8207158	Apr 10, 2023		U-1252	
		8207158	Apr 10, 2023		U-1253	
		8207158	Apr 10, 2023		U-1254	
		8207158	Apr 10, 2023		U-1255	
		8273734	Apr 10, 2023		U-1254	
		8273734	Apr 10, 2023		U-1255	
		8367657	Apr 10, 2023	DS DP	U-1252	
		8367657	Apr 10, 2023	DS DP	U-1253	
		8367657	Apr 10, 2023	DS DP	U-1254	
		8367657	Apr 10, 2023	DS DP	U-1255	
		8546379	Apr 10, 2023	DS DP	U-1252	
		8546379	Apr 10, 2023	DS DP	U-1253	
		8546379	Apr 10, 2023	DS DP	U-1254	
		8546379	Apr 10, 2023	DS DP	U-1255	
		8575149	Apr 10, 2023		U-1452	
		8697686	Dec 20, 2025	DS DP		
		8946207	Jun 16, 2024	DP		
		8980881	Dec 20, 2025		U-1252	
		8980881	Dec 20, 2025		U-1253	
		8980881	Dec 20, 2025		U-1254	
		8980881	Dec 20, 2025		U-1255	
		8999970	Feb 07, 2033		U-1688	
		8999970	Feb 07, 2033		U-1689	
		8999970	Feb 07, 2033		U-1692	
		9169213	Dec 06, 2032		U-1884	
		9169213	Dec 06, 2032		U-1885	
		9169213	Dec 06, 2032		U-1886	
		9169213	Dec 06, 2032		U-1887	
		9770455	Aug 31, 2031		U-2110	
<u>LORLATINIB - LORBRENA</u>						
N 210868	001	10420749	Jul 27, 2036	DS DP	U-2633	NCE Nov 02, 2023
		8680111	Mar 05, 2033	DS DP		ODE-217 Nov 02, 2025
						ODE-218 Nov 02, 2025
						ODE-219 Nov 02, 2025
<u>LORLATINIB - LORBRENA</u>						
N 210868	002	10420749	Jul 27, 2036	DS DP	U-2633	NCE Nov 02, 2023
		8680111	Mar 05, 2033	DS DP		ODE-217 Nov 02, 2025
						ODE-218 Nov 02, 2025
						ODE-219 Nov 02, 2025

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<u>LOTEPREDNOL ETABONATE - LOTEMAX</u>						
N 202872	001				M-229	Jul 20, 2021
<u>LOTEPREDNOL ETABONATE - LOTEMAX SM</u>						
N 208219	001				NS	Feb 22, 2022
<u>LOTEPREDNOL ETABONATE - INVELTYS</u>						
N 210565	001	10058511	May 03, 2033	DP U-2492	NP	Aug 22, 2021
		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>LOXAPINE - ADASUVE</u>						
N 022549	001	6716416	May 20, 2022	DP		
		7052679	Oct 26, 2021	DP		
		7078020	Oct 26, 2021	DP U-1375		
		7090830	Oct 26, 2021	DP		
		7458374	Aug 18, 2024	DP		
		7537009	Oct 28, 2024	DP		
		7585493	Oct 26, 2021	DP		
		7601337	Oct 26, 2021	DP		
		8074644	Jul 25, 2022	DP		
		8173107	Oct 26, 2021	DP		
		8235037	Oct 26, 2021	DP		
		8387612	Oct 23, 2026	DP		
		8955512	Oct 26, 2021	DP		
		8991387	May 21, 2024	DP		
		9370629	May 20, 2024	DP		
		9439907	Oct 26, 2021	DP		
		9440034	Oct 26, 2021	DP		
		9687487	Oct 26, 2021	DS DP		
<u>LUBIPROSTONE - AMITIZA</u>						
N 021908	001	6414016	Sep 05, 2020	U-1392	M-225	Apr 26, 2021
		6414016	Sep 05, 2020	U-717		
		6583174	Oct 16, 2020	DP		
		6982283	Dec 04, 2022	U-1391		
		7064148	Aug 30, 2022	U-1404		
		7064148	Aug 30, 2022	U-739		
		7417067	Oct 16, 2020	DP		
		8026393	Oct 25, 2027	DP		
		8071613	Sep 05, 2020	U-1203		
		8071613	Sep 05, 2020	U-1393		
		8088934	May 18, 2021	DS		
		8097649	Oct 16, 2020	DP		
		8097653	Nov 14, 2022	U-1214		
		8097653	Nov 14, 2022	U-1394		
		8114890	Sep 05, 2020	DP		
		8338639	Jan 23, 2027	DP		
		8389542	Nov 14, 2022	DP U-1345		
		8389542	Nov 14, 2022	DP U-1395		
		8748481	Sep 01, 2025	U-1520		
		8779187	Jan 23, 2027	DP		
<u>LUBIPROSTONE - AMITIZA</u>						
N 021908	002	6414016	Sep 05, 2020	U-874	M-225	Apr 26, 2021
		6583174	Oct 16, 2020	DP		
		7064148	Aug 30, 2022	U-739		
		7064148	Aug 30, 2022	U-873		
		7417067	Oct 16, 2020	DP		
		7795312	Sep 17, 2024	U-1085		
		8026393	Oct 25, 2027	DP		
		8071613	Sep 05, 2020	U-1202		
		8088934	May 18, 2021	DS		
		8097649	Oct 16, 2020	DP		
		8114890	Sep 05, 2020	DP		
		8338639	Jan 23, 2027	DP		
		8748481	Sep 01, 2025	U-1519		
		8779187	Jan 23, 2027	DP		

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<u>LUBIPROSTONE - AMITIZA</u>						
N 021908 002	6414016	Sep 05, 2020	U-874		M-225	Apr 26, 2021
	6583174	Oct 16, 2020	DP			
	7064148	Aug 30, 2022	U-739			
	7064148	Aug 30, 2022	U-873			
	7417067	Oct 16, 2020	DP			
	7795312	Sep 17, 2024	U-1085			
	8026393	Oct 25, 2027	DP			
	8071613	Sep 05, 2020	U-1202			
	8088934	May 18, 2021	DS			
	8097649	Oct 16, 2020	DP			
	8114890	Sep 05, 2020	DP			
	8338639	Jan 23, 2027	DP			
	8748481	Sep 01, 2025	U-1519			
	8779187	Jan 23, 2027	DP			
<u>LULICONAZOLE - LUZU</u>						
N 204153 001	5900488	Jan 18, 2020	DS DP		NPP	Feb 20, 2021
	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>LUMATEPERONE TOSYLATE - CAPLYTA</u>						
N 209500 001	10464938	Mar 12, 2028	DP		NCE	Dec 20, 2024
	7183282	Jun 15, 2020	DS DP			
	8598119	Dec 28, 2029	U-543			
	8648077	Dec 01, 2029	DS DP			
	9199995	Mar 12, 2029	U-2713			
	9586960	Mar 12, 2029	DS DP			
	9616061	May 27, 2029	DP			
	9956227	Dec 03, 2034	U-2714			
	RE39680	Jun 15, 2020	DS DP U-543			
<u>LURASIDONE HYDROCHLORIDE - LATUDA</u>						
N 200603 001	8729085	May 26, 2026	DP		M-195	Jan 27, 2020
	8729085*PED	Nov 26, 2026			NPP	Jan 27, 2020
	8883794	May 26, 2026	DP		NPP	Mar 05, 2021
	8883794*PED	Nov 26, 2026			PED	Jul 27, 2020
	9174975	Feb 20, 2024	U-1770		PED	Jul 27, 2020
	9174975*PED	Aug 20, 2024				
	9259423	May 23, 2031	U-1822			
	9259423*PED	Nov 23, 2031				
	9555027	May 26, 2026	DP U-543			
	9815827	Feb 20, 2024	U-2166			
	9815827	Feb 20, 2024	U-543			
	9827242	May 23, 2031	U-2199			
	9827242	May 23, 2031	U-2201			
	9907794	May 26, 2026	DP			
	RE45573	Jun 23, 2025	DS			
	RE45573*PED	Dec 23, 2025				
<u>LURASIDONE HYDROCHLORIDE - LATUDA</u>						
N 200603 002	8729085	May 26, 2026	DP		NPP	Jan 27, 2020
	8729085*PED	Nov 26, 2026			NPP	Mar 05, 2021
	8883794	May 26, 2026	DP		PED	Jul 27, 2020
	8883794*PED	Nov 26, 2026				
	9174975	Feb 20, 2024	U-1770			
	9174975*PED	Aug 20, 2024				
	9259423	May 23, 2031	U-1822			
	9259423*PED	Nov 23, 2031				
	9555027	May 26, 2026	DP U-543			
	9815827	Feb 20, 2024	U-2166			
	9815827	Feb 20, 2024	U-543			
	9827242	May 23, 2031	U-2199			
	9827242	May 23, 2031	U-2201			
	9907794	May 26, 2026	DP			
	RE45573	Jun 23, 2025	DS			
	RE45573*PED	Dec 23, 2025				

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<u>LURASIDONE HYDROCHLORIDE - LATUDA</u>						
N 200603 003	8729085	May 26, 2026	DP		M-195	Jan 27, 2020
	8729085*PED	Nov 26, 2026			NPP	Jan 27, 2020
	8883794	May 26, 2026	DP		NPP	Mar 05, 2021
	8883794*PED	Nov 26, 2026			PED	Jul 27, 2020
	9174975	Feb 20, 2024	U-1770		PED	Jul 27, 2020
	9174975*PED	Aug 20, 2024				
	9259423	May 23, 2031	U-1822			
	9259423*PED	Nov 23, 2031				
	9555027	May 26, 2026	DP U-543			
	9815827	Feb 20, 2024	U-2166			
	9815827	Feb 20, 2024	U-543			
	9827242	May 23, 2031	U-2199			
	9827242	May 23, 2031	U-2201			
	9907794	May 26, 2026	DP			
	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				
	9174975	Feb 20, 2024	U-1770			
	9174975*PED	Aug 20, 2024				
	9259423	May 23, 2031	U-1822			
	9259423*PED	Nov 23, 2031				
	9555027	May 26, 2026	DP U-543			
	9815827	Feb 20, 2024	U-2166			
	9815827	Feb 20, 2024	U-543			
	9827242	May 23, 2031	U-2199			
	9827242	May 23, 2031	U-2201			
	9907794	May 26, 2026	DP			
	RE45573	Jun 23, 2025	DS			
	RE45573*PED	Dec 23, 2025				
<u>LURASIDONE HYDROCHLORIDE - LATUDA</u>						
N 200603 005	8729085	May 26, 2026	DP		M-195	Jan 27, 2020
	8729085*PED	Nov 26, 2026			NPP	Jan 27, 2020
	8883794	May 26, 2026	DP		NPP	Mar 05, 2021
	8883794*PED	Nov 26, 2026			PED	Jul 27, 2020
	9174975	Feb 20, 2024	U-1770		PED	Jul 27, 2020
	9174975*PED	Aug 20, 2024				
	9259423	May 23, 2031	U-1822			
	9259423*PED	Nov 23, 2031				
	9555027	May 26, 2026	DP U-543			
	9815827	Feb 20, 2024	U-2166			
	9815827	Feb 20, 2024	U-543			
	9827242	May 23, 2031	U-2199			
	9827242	May 23, 2031	U-2201			
	9907794	May 26, 2026	DP			
	RE45573	Jun 23, 2025	DS			
	RE45573*PED	Dec 23, 2025				
<u>LUSUTROMBOPAG - MULPLETA</u>						
N 210923 001	7601746	Sep 05, 2024	DS DP U-2344		NCE	Jul 31, 2023
	8530668	Jan 21, 2030	DS DP			
	8889722	Jul 29, 2028	DS DP			
	9427402	Sep 29, 2031	DP			
<u>LUTETIUM DOTATATE LU-177 - LUTATHERA</u>						
N 208700 001					NCE	Jan 26, 2023
					ODE-166	Jan 26, 2025
<u>MACIMORELIN ACETATE - MACRILEN</u>						
N 205598 001	6861409	Aug 01, 2022	DS DP U-2220		NCE	Dec 20, 2022
	8192719	Oct 12, 2027	U-2220		ODE-170	Dec 20, 2024

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<u>MACITENTAN - OPSUMIT</u>						
N 204410 001	7094781	Dec 05, 2025	DS DP		ODE-54	Oct 18, 2020
	8268847	Apr 18, 2029		U-1446		
	8367685	Oct 04, 2028		DP U-1445		
	9265762	May 29, 2027		DP U-1820		
<u>MAGNESIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE - NORMOCARB HF 25</u>						
N 021910 001	7300674	Mar 04, 2023	DP U-785			
<u>MAGNESIUM SULFATE; POTASSIUM SULFATE; SODIUM SULFATE - SUPREP BOWEL PREP KIT</u>						
N 022372 001	6946149	Mar 07, 2023	DP U-837			
<u>MALATHION - OVIDE</u>						
N 018613 001	7560445	Feb 01, 2027	DS DP U-986			
	7977324	Aug 14, 2026	DP			
<u>MARAVIROC - SELZENTRY</u>						
N 022128 001	6667314	Aug 06, 2021	DS DP U-824			
	7368460	Nov 25, 2022		U-824		
	7576097	May 25, 2021	DS			
<u>MARAVIROC - SELZENTRY</u>						
N 022128 002	6667314	Aug 06, 2021	DS DP U-824			
	7368460	Nov 25, 2022		U-824		
	7576097	May 25, 2021	DS			
<u>MARAVIROC - SELZENTRY</u>						
N 022128 003	6667314	Aug 06, 2021	DS DP U-824			
	7368460	Nov 25, 2022		U-824		
	7576097	May 25, 2021	DS			
<u>MARAVIROC - SELZENTRY</u>						
N 022128 004	6667314	Aug 06, 2021	DS DP U-824			
	7368460	Nov 25, 2022		U-824		
	7576097	May 25, 2021	DS			
<u>MARAVIROC - SELZENTRY</u>						
N 208984 001	6667314	Aug 06, 2021	DS DP U-824			
	7368460	Nov 25, 2022		U-824		
	7576097	May 25, 2021	DS			
<u>MECHLORETHAMINE HYDROCHLORIDE - VALCHLOR</u>						
N 202317 001	7838564	Mar 07, 2026	DP		ODE-51	Aug 23, 2020
	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>MEDROXYPROGESTERONE ACETATE - DEPO-SUBO PROVERA 104</u>						
N 021583 001	6495534	May 15, 2020	DP			
<u>MEGESTROL ACETATE - MEGACE ES</u>						
N 021778 001	6592903	Sep 21, 2020	DP			
	7101576	Apr 22, 2024		U-755		
	9040088	Apr 22, 2024		U-755		
	9101540	Apr 22, 2024	DP U-755			
	9101549	Apr 22, 2024		U-755		
	9107827	Apr 22, 2024		U-755		
<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		

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<u>MELOXICAM - ANJESO</u>						
N 210583	001	10463673	Feb 24, 2024	DP U-2750	NP	Feb 20, 2023
		10471067	Feb 24, 2024	DP U-2750		
		8512727	Dec 25, 2022	DP U-2750		
		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 - EVOMELA</u>						
N 207155	001	10040872	Jan 30, 2034	DP	ODE-110	Mar 10, 2023
		8410077	Mar 13, 2029	DP		
		9200088	Mar 13, 2029	DP		
		9493582	Feb 27, 2033	DP		
<u>MEMANTINE HYDROCHLORIDE - NAMENDA XR</u>						
N 022525	001	8039009	Mar 24, 2029	U-539		
		8039009*PED	Sep 24, 2029			
		8168209	Nov 22, 2025	DP		
		8168209*PED	May 22, 2026			
		8173708	Nov 22, 2025	U-539		
		8173708*PED	May 22, 2026			
		8283379	Nov 22, 2025	U-539		
		8283379*PED	May 22, 2026			
		8329752	Nov 22, 2025	DP		
		8329752*PED	May 22, 2026			
		8362085	Nov 22, 2025	U-539		
		8362085*PED	May 22, 2026			
<u>MEMANTINE HYDROCHLORIDE - NAMENDA XR</u>						
N 022525	002	8039009	Mar 24, 2029	U-539		
		8039009*PED	Sep 24, 2029			
		8168209	Nov 22, 2025	DP		
		8168209*PED	May 22, 2026			
		8173708	Nov 22, 2025	U-539		
		8173708*PED	May 22, 2026			
		8283379	Nov 22, 2025	U-539		
		8283379*PED	May 22, 2026			
		8329752	Nov 22, 2025	DP		
		8329752*PED	May 22, 2026			
		8362085	Nov 22, 2025	U-539		
		8362085*PED	May 22, 2026			
<u>MEMANTINE HYDROCHLORIDE - NAMENDA XR</u>						
N 022525	003	8039009	Mar 24, 2029	U-539		
		8039009*PED	Sep 24, 2029			
		8168209	Nov 22, 2025	DP		
		8168209*PED	May 22, 2026			
		8173708	Nov 22, 2025	U-539		
		8173708*PED	May 22, 2026			
		8283379	Nov 22, 2025	U-539		
		8283379*PED	May 22, 2026			
		8329752	Nov 22, 2025	DP		
		8329752*PED	May 22, 2026			
		8362085	Nov 22, 2025	U-539		
		8362085*PED	May 22, 2026			
<u>MEMANTINE HYDROCHLORIDE - NAMENDA XR</u>						
N 022525	004	8039009	Mar 24, 2029	U-539		
		8039009*PED	Sep 24, 2029			
		8168209	Nov 22, 2025	DP		
		8168209*PED	May 22, 2026			
		8173708	Nov 22, 2025	U-539		
		8173708*PED	May 22, 2026			
		8283379	Nov 22, 2025	U-539		
		8283379*PED	May 22, 2026			
		8329752	Nov 22, 2025	DP		
		8329752*PED	May 22, 2026			

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<u>MEMANTINE HYDROCHLORIDE - NAMENDA XR</u>						
N 022525	004	8362085	Nov 22, 2025		U-539	
		8362085*PED	May 22, 2026			
		8598233	Nov 22, 2025	DP		
		8598233*PED	May 22, 2026			
<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>MEQUINOL; TRETINOIN - SOLAGE</u>						
N 020922	001	6353029	Aug 24, 2020			
<u>MERCAPTOPYRINE - PURIXAN</u>						
N 205919	001				ODE-65	Apr 28, 2021
<u>MEROPENEM; VABORBACTAM - VABOMERE</u>						
N 209776	001	10172874	Aug 08, 2031	DP		
		10183034	Aug 08, 2031		U-2490	
		8680136	Aug 17, 2031	DS DP		
		9694025	Aug 08, 2031		U-2120	
<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 - ASACOL HD</u>						
N 021830	001	6893662	Nov 15, 2021	DP	U-141	
		8580302	Nov 15, 2021	DP		
		9089492	Nov 15, 2021	DP		
<u>MESALAMINE - LIALDA</u>						
N 022000	001	6773720	Jun 08, 2020	DP		
<u>MESALAMINE - APRISO</u>						
N 022301	001	8865688	May 01, 2030		U-1310	
<u>MESALAMINE - DELZICOL</u>						
N 204412	001	6649180	Apr 13, 2020	DP		
<u>METAXALONE - SKELAXIN</u>						
N 013217	003	7122566	Feb 06, 2026		U-915	
		7714006	Dec 03, 2021		U-1050	
<u>METFORMIN HYDROCHLORIDE - FORTAMET</u>						
N 021574	001	6790459	Mar 17, 2021		U-604	
		6866866	Mar 17, 2021	DP		
<u>METFORMIN HYDROCHLORIDE - FORTAMET</u>						
N 021574	002	6790459	Mar 17, 2021		U-604	
		6866866	Mar 17, 2021	DP		
<u>METFORMIN HYDROCHLORIDE - RIOMET</u>						
N 021591	001	6890957	Aug 07, 2021	DP		
<u>METFORMIN HYDROCHLORIDE - GLUMETZA</u>						
N 021748	001	6488962	Jun 20, 2020	DS DP		
		6723340	Oct 25, 2021	DS DP		
<u>METFORMIN HYDROCHLORIDE - GLUMETZA</u>						
N 021748	002	7780987	Mar 23, 2025	DS DP		
		8323692	Mar 30, 2023	DP		

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<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 6790459	Mar 17, 2021		U-974		
	6866866	Mar 17, 2021	DP			
	7785627	Jul 31, 2026	DP			
	7959946	Jul 31, 2026	DP			
	8470368	Sep 19, 2023	DP			
	8668931	Sep 19, 2023	DP			
	9060941	Sep 19, 2023	DP			
<u>METFORMIN HYDROCHLORIDE; PIOGLITAZONE HYDROCHLORIDE - ACTOPLUS MET XR</u>						
N 022024	002 6790459	Mar 17, 2021		U-974		
	6866866	Mar 17, 2021	DP			
	7785627	Jul 31, 2026	DP			
	7959946	Jul 31, 2026	DP			
	8470368	Sep 19, 2023	DP			
	8668931	Sep 19, 2023	DP			
	9060941	Sep 19, 2023	DP			
<u>METFORMIN HYDROCHLORIDE; ROSIGLITAZONE MALEATE - AVANDAMET</u>						
N 021410	001 8236345	Oct 07, 2022	DP			
<u>METFORMIN HYDROCHLORIDE; ROSIGLITAZONE MALEATE - AVANDAMET</u>						
N 021410	002 8236345	Oct 07, 2022	DP			
<u>METFORMIN HYDROCHLORIDE; ROSIGLITAZONE MALEATE - AVANDAMET</u>						
N 021410	003 8236345	Oct 07, 2022	DP			
<u>METFORMIN HYDROCHLORIDE; ROSIGLITAZONE MALEATE - AVANDAMET</u>						
N 021410	004 8236345	Oct 07, 2022	DP			
<u>METFORMIN HYDROCHLORIDE; ROSIGLITAZONE MALEATE - AVANDAMET</u>						
N 021410	005 8236345	Oct 07, 2022	DP			
<u>METFORMIN HYDROCHLORIDE; SAXAGLIPTIN HYDROCHLORIDE - KOMBIGLYZE XR</u>						
N 200678	001 8628799	Jul 13, 2025	DP		M-198	Feb 27, 2020
	9339472	Jul 13, 2025	DP			
	RE44186	Jul 31, 2023	DS DP U-1097			
	RE44186	Jul 31, 2023	DS DP U-1838			
<u>METFORMIN HYDROCHLORIDE; SAXAGLIPTIN HYDROCHLORIDE - KOMBIGLYZE XR</u>						
N 200678	002 8628799	Jul 13, 2025	DP		M-198	Feb 27, 2020
	9339472	Jul 13, 2025	DP			
	RE44186	Jul 31, 2023	DS DP U-1097			
	RE44186	Jul 31, 2023	DS DP U-1838			
<u>METFORMIN HYDROCHLORIDE; SAXAGLIPTIN HYDROCHLORIDE - KOMBIGLYZE XR</u>						
N 200678	003 8628799	Jul 13, 2025	DP		M-198	Feb 27, 2020
	9339472	Jul 13, 2025	DP			
	RE44186	Jul 31, 2023	DS DP U-1097			
	RE44186	Jul 31, 2023	DS DP U-1838			
<u>METFORMIN HYDROCHLORIDE; SITAGLIPTIN PHOSPHATE - JANUMET</u>						
N 022044	001 6699871	Jul 26, 2022	DS DP U-802			
	7125873	Jul 26, 2022	DP U-1036			
	7125873	Jul 26, 2022	DP U-1038			
	7125873	Jul 26, 2022	DP U-803			
	7326708	Nov 24, 2026	DS DP U-802			
	8414921	Jul 21, 2028	DP U-1036			

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<u>METFORMIN HYDROCHLORIDE; SITAGLIPTIN PHOSPHATE - JANUMET</u>						
N 022044	002	6699871	Jul 26, 2022	DS DP	U-802	
		7125873	Jul 26, 2022		DP U-1036	
		7125873	Jul 26, 2022		DP U-1038	
		7125873	Jul 26, 2022		DP U-803	
		7326708	Nov 24, 2026	DS DP	U-802	
		8414921	Jul 21, 2028		DP U-1036	
<u>METFORMIN HYDROCHLORIDE; SITAGLIPTIN PHOSPHATE - JANUMET XR</u>						
N 202270	001	6699871	Jul 26, 2022	DS DP	U-1227	
		7125873	Jul 26, 2022		DP U-1227	
		7326708	Nov 24, 2026	DS DP	U-1227	
<u>METFORMIN HYDROCHLORIDE; SITAGLIPTIN PHOSPHATE - JANUMET XR</u>						
N 202270	002	6699871	Jul 26, 2022	DS DP	U-1227	
		7125873	Jul 26, 2022		DP U-1227	
		7326708	Nov 24, 2026	DS DP	U-1227	
<u>METFORMIN HYDROCHLORIDE; SITAGLIPTIN PHOSPHATE - JANUMET XR</u>						
N 202270	003	6699871	Jul 26, 2022	DS DP	U-1227	
		7125873	Jul 26, 2022		DP U-1227	
		7326708	Nov 24, 2026	DS DP	U-1227	
<u>METHOTREXATE - OTREXUP</u>						
N 204824	001	6746429	Apr 12, 2020		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	
		8945063	Mar 19, 2030		DP U-1442	
		9421333	Mar 19, 2030		DP U-1442	
		9533102	Jan 24, 2026		DP	
		9629959	Jan 24, 2026		DP	
<u>METHOTREXATE - OTREXUP</u>						
N 204824	002	6746429	Apr 12, 2020		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	
		8945063	Mar 19, 2030		DP U-1442	
		9421333	Mar 19, 2030		DP U-1442	
		9533102	Jan 24, 2026		DP	
		9629959	Jan 24, 2026		DP	
<u>METHOTREXATE - OTREXUP</u>						
N 204824	003	6746429	Apr 12, 2020		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	
		8945063	Mar 19, 2030		DP U-1442	
		9421333	Mar 19, 2030		DP U-1442	
		9533102	Jan 24, 2026		DP	
		9629959	Jan 24, 2026		DP	
<u>METHOTREXATE - OTREXUP</u>						
N 204824	004	6746429	Apr 12, 2020		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	
		8945063	Mar 19, 2030		DP U-1442	
		9421333	Mar 19, 2030		DP U-1442	
		9533102	Jan 24, 2026		DP	
		9629959	Jan 24, 2026		DP	
<u>METHOTREXATE - OTREXUP</u>						
N 204824	005	6746429	Apr 12, 2020		DP	
		8021335	Oct 04, 2026		DP	
		8480631	Mar 19, 2030		DP U-1442	

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<u>METHOTREXATE - OTREXUP</u>						
N 204824	005	8562564	Jan 24, 2026	DP		
		8579865	Mar 19, 2030	DP U-1442		
		8945063	Mar 19, 2030	DP U-1442		
		9421333	Mar 19, 2030	DP U-1442		
		9533102	Jan 24, 2026	DP		
		9629959	Jan 24, 2026	DP		
<u>METHOTREXATE - OTREXUP</u>						
N 204824	006	6746429	Apr 12, 2020	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		
		8945063	Mar 19, 2030	DP U-1442		
		9421333	Mar 19, 2030	DP U-1442		
		9533102	Jan 24, 2026	DP		
		9629959	Jan 24, 2026	DP		
<u>METHOTREXATE - OTREXUP</u>						
N 204824	007	6746429	Apr 12, 2020	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		
		8945063	Mar 19, 2030	DP U-1442		
		9421333	Mar 19, 2030	DP U-1442		
		9533102	Jan 24, 2026	DP		
		9629959	Jan 24, 2026	DP		
<u>METHOTREXATE - OTREXUP</u>						
N 204824	008	6746429	Apr 12, 2020	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		
		8945063	Mar 19, 2030	DP U-1442		
		9421333	Mar 19, 2030	DP U-1442		
		9533102	Jan 24, 2026	DP		
		9629959	Jan 24, 2026	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		
<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		

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<u>METHOTREXATE SODIUM - XATMEP</u>						
N 208400 001	10231927	Jan 02, 2033		U-1349	ODE-137	Apr 25, 2024
	10231927	Jan 02, 2033		U-1699	ODE-138	Apr 25, 2024
	9259427	Jan 02, 2033	DP			
	9855215	Jan 02, 2033	DP			
<u>METHYLENE BLUE - PROVAYBLUE</u>						
N 204630 001					ODE-113	Apr 08, 2023
<u>METHYLNALTREXONE BROMIDE - RELISTOR</u>						
N 021964 001	10376584	Apr 08, 2024	DP	U-1185		
	8247425	Dec 31, 2030		U-1185		
	8420663	Sep 30, 2029		U-1185		
	8552025	Apr 08, 2024	DP			
	8822490	Sep 30, 2029	DP	U-1185		
	9180125	Sep 30, 2029	DP	U-1185		
	9492445	Sep 30, 2029	DP	U-1185		
	9669096	Apr 08, 2024	DP			
<u>METHYLNALTREXONE BROMIDE - RELISTOR</u>						
N 021964 002	10376584	Apr 08, 2024	DP	U-1185		
	8247425	Dec 31, 2030		U-1185		
	8420663	Sep 30, 2029		U-1185		
	8552025	Apr 08, 2024	DP			
	8822490	Sep 30, 2029	DP	U-1185		
	9180125	Sep 30, 2029	DP	U-1185		
	9492445	Sep 30, 2029	DP	U-1185		
	9669096	Apr 08, 2024	DP			
<u>METHYLNALTREXONE BROMIDE - RELISTOR</u>						
N 021964 003	10376584	Apr 08, 2024	DP	U-1185		
	8247425	Dec 31, 2030		U-1185		
	8420663	Sep 30, 2029		U-1185		
	8552025	Apr 08, 2024	DP			
	8822490	Sep 30, 2029	DP	U-1185		
	9180125	Sep 30, 2029	DP	U-1185		
	9492445	Sep 30, 2029	DP	U-1185		
	9669096	Apr 08, 2024	DP			
<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		

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<u>METHYLPHENIDATE - COTEMPLA XR-ODT</u>						
N 205489 001	8840924	Jun 05, 2026	DP		NP	Jun 19, 2020
	9072680	Jun 28, 2032	DP			
	9089496	Jun 28, 2032	DP			
<u>METHYLPHENIDATE - COTEMPLA XR-ODT</u>						
N 205489 002	8840924	Jun 05, 2026	DP		NP	Jun 19, 2020
	9072680	Jun 28, 2032	DP			
	9089496	Jun 28, 2032	DP			
<u>METHYLPHENIDATE - COTEMPLA XR-ODT</u>						
N 205489 003	8840924	Jun 05, 2026	DP		NP	Jun 19, 2020
	9072680	Jun 28, 2032	DP			
	9089496	Jun 28, 2032	DP			
<u>METHYLPHENIDATE HYDROCHLORIDE - METADATE CD</u>						
N 021259 001	6344215	Oct 27, 2020	DP			
<u>METHYLPHENIDATE HYDROCHLORIDE - METADATE CD</u>						
N 021259 002	6344215	Oct 27, 2020	DP			
<u>METHYLPHENIDATE HYDROCHLORIDE - METADATE CD</u>						
N 021259 003	6344215	Oct 27, 2020	DP			
<u>METHYLPHENIDATE HYDROCHLORIDE - METADATE CD</u>						
N 021259 004	6344215	Oct 27, 2020	DP			
<u>METHYLPHENIDATE HYDROCHLORIDE - METHYLIN</u>						
N 021419 001	7691880	Oct 07, 2024	DP			
<u>METHYLPHENIDATE HYDROCHLORIDE - METHYLIN</u>						
N 021419 002	7691880	Oct 07, 2024	DP			
<u>METHYLPHENIDATE HYDROCHLORIDE - OUILIVANT 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 - APTENSIO XR</u>						
N 205831 001	10039719*PED	Jun 16, 2020				
	6419960*PED	Jun 16, 2020				
	7083808*PED	Jun 16, 2020				
	7247318*PED	Jun 16, 2020				
	7438930*PED	Jun 16, 2020				
	8580310*PED	Jun 16, 2020				
	9066869*PED	Jun 16, 2020				
	9801823*PED	Jun 16, 2020				
<u>METHYLPHENIDATE HYDROCHLORIDE - APTENSIO XR</u>						
N 205831 002	10039719*PED	Jun 16, 2020				
	6419960*PED	Jun 16, 2020				
	7083808*PED	Jun 16, 2020				
	7247318*PED	Jun 16, 2020				
	7438930*PED	Jun 16, 2020				
	8580310*PED	Jun 16, 2020				
	9066869*PED	Jun 16, 2020				
	9801823*PED	Jun 16, 2020				
<u>METHYLPHENIDATE HYDROCHLORIDE - APTENSIO XR</u>						
N 205831 003	10039719*PED	Jun 16, 2020				
	6419960*PED	Jun 16, 2020				
	7083808*PED	Jun 16, 2020				
	7247318*PED	Jun 16, 2020				
	7438930*PED	Jun 16, 2020				
	8580310*PED	Jun 16, 2020				
	9066869*PED	Jun 16, 2020				
	9801823*PED	Jun 16, 2020				

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APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>METHYLPHENIDATE HYDROCHLORIDE - APTENSIO XR</u>						
N 205831 004	10039719*PED	Jun 16, 2020				
	6419960*PED	Jun 16, 2020				
	7083808*PED	Jun 16, 2020				
	7247318*PED	Jun 16, 2020				
	7438930*PED	Jun 16, 2020				
	8580310*PED	Jun 16, 2020				
	9066869*PED	Jun 16, 2020				
	9801823*PED	Jun 16, 2020				
<u>METHYLPHENIDATE HYDROCHLORIDE - APTENSIO XR</u>						
N 205831 005	10039719*PED	Jun 16, 2020				
	6419960*PED	Jun 16, 2020				
	7083808*PED	Jun 16, 2020				
	7247318*PED	Jun 16, 2020				
	7438930*PED	Jun 16, 2020				
	8580310*PED	Jun 16, 2020				
	9066869*PED	Jun 16, 2020				
	9801823*PED	Jun 16, 2020				
<u>METHYLPHENIDATE HYDROCHLORIDE - APTENSIO XR</u>						
N 205831 006	10039719*PED	Jun 16, 2020				
	6419960*PED	Jun 16, 2020				
	7083808*PED	Jun 16, 2020				
	7247318*PED	Jun 16, 2020				
	7438930*PED	Jun 16, 2020				
	8580310*PED	Jun 16, 2020				
	9066869*PED	Jun 16, 2020				
	9801823*PED	Jun 16, 2020				
<u>METHYLPHENIDATE HYDROCHLORIDE - APTENSIO XR</u>						
N 205831 007	10039719*PED	Jun 16, 2020				
	6419960*PED	Jun 16, 2020				
	7083808*PED	Jun 16, 2020				
	7247318*PED	Jun 16, 2020				
	7438930*PED	Jun 16, 2020				
	8580310*PED	Jun 16, 2020				
	9066869*PED	Jun 16, 2020				
	9801823*PED	Jun 16, 2020				
<u>METHYLPHENIDATE HYDROCHLORIDE - QUILLICHEW ER</u>						
N 207960 001	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	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	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	NP	Aug 08, 2021
	10292937	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		

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<u>METHYLPHENIDATE HYDROCHLORIDE - JORNAY PM</u>						
N 209311 001	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 002	10182995	Mar 23, 2032	DP		NP	Aug 08, 2021
	10292937	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 003	10182995	Mar 23, 2032	DP		NP	Aug 08, 2021
	10292937	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 004	10182995	Mar 23, 2032	DP		NP	Aug 08, 2021
	10292937	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		NP	Aug 08, 2021
	10292937	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		NP	Feb 27, 2022
	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		
	9974752	Oct 30, 2035	DP			
<u>METHYLPHENIDATE HYDROCHLORIDE - ADHANSIA XR</u>						
N 212038 002	10111839	Oct 30, 2035	U-2357		NP	Feb 27, 2022
	10292938	Oct 30, 2035	DP			
	10292939	Oct 30, 2035	DP	U-2357		
	10449159	Oct 30, 2035	U-2357			

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<u>METHYLPHENIDATE HYDROCHLORIDE - ADHANSIA XR</u>						
N 212038 002	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			
	9974752	Oct 30, 2035	DP			
<u>METHYLPHENIDATE HYDROCHLORIDE - ADHANSIA XR</u>						
N 212038 003	10111839	Oct 30, 2035	U-2357		NP	Feb 27, 2022
	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			
	9974752	Oct 30, 2035	DP			
<u>METHYLPHENIDATE HYDROCHLORIDE - ADHANSIA XR</u>						
N 212038 004	10111839	Oct 30, 2035	U-2357		NP	Feb 27, 2022
	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			
	9974752	Oct 30, 2035	DP			
<u>METHYLPHENIDATE HYDROCHLORIDE - ADHANSIA XR</u>						
N 212038 005	10111839	Oct 30, 2035	U-2357		NP	Feb 27, 2022
	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			
	9974752	Oct 30, 2035	DP			
<u>METHYLPHENIDATE HYDROCHLORIDE - ADHANSIA XR</u>						
N 212038 006	10111839	Oct 30, 2035	U-2357		NP	Feb 27, 2022
	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			
	9974752	Oct 30, 2035	DP			
<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			

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<u>METOPROLOL SUCCINATE - KAPSPARGO SPRINKLE</u>						
N 210428	004 9504655	Jul 09, 2035	DP			
	9700530	Jul 09, 2035	DP			
<u>METRONIDAZOLE - METROGEL</u>						
N 021789	001 6881726	Feb 21, 2022	DP U-743			
	7348317	Feb 21, 2022	DP U-743			
<u>METRONIDAZOLE - VANDAZOLE</u>						
N 021806	001 7456207	Sep 22, 2024	DP			
<u>METRONIDAZOLE - NUVESSA</u>						
N 205223	001 10238634	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>MICAFUNGIN SODIUM - MYCAMINE</u>						
N 021506	002 6774104	Jan 08, 2021	DP U-650		I-821	Dec 20, 2022
	6774104	Jan 08, 2021	DP U-845		PED	Jun 20, 2023
	6774104*PED	Jul 08, 2021				
<u>MICAFUNGIN SODIUM - MYCAMINE</u>						
N 021506	003 6774104	Jan 08, 2021	DP U-650		I-821	Dec 20, 2022
	6774104	Jan 08, 2021	DP U-845		PED	Jun 20, 2023
	6774104*PED	Jul 08, 2021				
<u>MICONAZOLE - ORAVIG</u>						
N 022404	001 6916485	Sep 11, 2022	DP U-1051			
	7651698	Sep 11, 2022	U-1051			
	8518442	Sep 11, 2022	DP			
<u>MICONAZOLE NITRATE; MICONAZOLE NITRATE - MONISTAT 1 COMBINATION PACK</u>						
N 021308	001 6153635	Nov 28, 2020				Y
<u>MICONAZOLE NITRATE; PETROLATUM, WHITE; 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		NP	May 21, 2022
	8809322	Jan 18, 2028	DP		ODE-243	May 17, 2026
	9289432	Jan 18, 2028	DP U-2526			
	9687495	Jan 18, 2028	DP U-2526			
<u>MIDAZOLAM HYDROCHLORIDE - SEIZALAM</u>						
N 209566	001				ODE-207	Sep 14, 2025
<u>MIDOSTAURIN - RYDAPT</u>						
N 207997	001 7973031	Oct 17, 2024	U-2007		NCE	Apr 28, 2022
	8222244	Oct 29, 2022	U-2007		ODE-140	Apr 28, 2024
	8575146	Dec 02, 2030	U-2008		ODE-141	Apr 28, 2024
<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			
	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		NCE	Aug 10, 2023
	10251873	May 30, 2038	U-2371		ODE-205	Aug 10, 2025
	10383864	May 16, 2027	U-2371			

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<u>MIGALASTAT HYDROCHLORIDE - GALAFOLD</u>						
N 208623	001	10406143	May 16, 2027			
		10471053	May 30, 2038			
		10525045	Apr 28, 2028			
		8592362	Feb 12, 2029			
		9000011	May 16, 2027			
		9095584	Feb 12, 2029			
		9480682	May 16, 2027			
		9987263	May 16, 2027			
		9999618	Apr 28, 2028			
		9999618	Apr 28, 2028			
<u>MILNACIPRAN HYDROCHLORIDE - SAVELLA</u>						
N 022256	001	6602911	Jan 14, 2023			
		6992110	Nov 05, 2021			
		7888342	Nov 05, 2021			
		7994220	Sep 19, 2029			
<u>MILNACIPRAN HYDROCHLORIDE - SAVELLA</u>						
N 022256	002	6602911	Jan 14, 2023			
		6992110	Nov 05, 2021			
		7888342	Nov 05, 2021			
		7994220	Sep 19, 2029			
<u>MILNACIPRAN HYDROCHLORIDE - SAVELLA</u>						
N 022256	003	6602911	Jan 14, 2023			
		6992110	Nov 05, 2021			
		7888342	Nov 05, 2021			
		7994220	Sep 19, 2029			
<u>MILNACIPRAN HYDROCHLORIDE - SAVELLA</u>						
N 022256	004	6602911	Jan 14, 2023			
		6992110	Nov 05, 2021			
		7888342	Nov 05, 2021			
		7994220	Sep 19, 2029			
<u>MILTEFOSINE - IMPAVIDO</u>						
N 204684	001				ODE-63	Mar 19, 2021
<u>MINOCYCLINE HYDROCHLORIDE - MINOCIN</u>						
N 050444	001	9084802	May 12, 2031			
		9278105	May 12, 2031			
<u>MINOCYCLINE HYDROCHLORIDE - ARESTIN</u>						
N 050781	001	6682348	Mar 29, 2022	DP		
		7699609	Mar 29, 2022	DP		
<u>MINOCYCLINE HYDROCHLORIDE - SOLODYN</u>						
N 050808	001	7790705	Jun 24, 2025			
		7919483	Mar 07, 2027			
		8252776	Jun 24, 2025			
		8268804	Jun 24, 2025			
<u>MINOCYCLINE HYDROCHLORIDE - SOLODYN</u>						
N 050808	002	7541347	Apr 02, 2027			
		7544373	Apr 02, 2027	DP		
		7790705	Jun 24, 2025			
		7919483	Mar 07, 2027			
		8252776	Jun 24, 2025			
		8268804	Jun 24, 2025			
<u>MINOCYCLINE HYDROCHLORIDE - SOLODYN</u>						
N 050808	003	7790705	Jun 24, 2025			
		7919483	Mar 07, 2027			
		8252776	Jun 24, 2025			
		8268804	Jun 24, 2025			
<u>MINOCYCLINE HYDROCHLORIDE - SOLODYN</u>						
N 050808	004	7790705	Jun 24, 2025			
		7919483	Mar 07, 2027			
		8252776	Jun 24, 2025			

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<u>MINOCYCLINE HYDROCHLORIDE - SOLODYN</u>						
N 050808	004	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
		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 - 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
		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>MIPOMERSEN SODIUM - KYNAMRO</u>						
N 203568	001	7015315	Mar 21, 2023	DS	ODE-41	Jan 29, 2020
		7101993	Sep 05, 2023	DS		

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<u>MIPOMERSEN SODIUM - KYNAMRO</u>						
N 203568	001	7407943	Aug 01, 2021	U-1353		
		7511131	Jan 29, 2027	DS		
<u>MIRABEGRON - MYRBETRIO</u>						
N 202611	001	6346532	Mar 27, 2022	DS DP	I-777	Apr 27, 2021
		6562375	Aug 01, 2020	DP		
		7342117	Nov 04, 2023	DS		
		7982049	Nov 04, 2023	DP		
		8772315	Oct 30, 2028	U-2300		
		8835474	Nov 04, 2023	U-1527		
		RE44872	Nov 04, 2023	U-1527		
<u>MIRABEGRON - MYRBETRIO</u>						
N 202611	002	6346532	Mar 27, 2022	DS DP	I-777	Apr 27, 2021
		6562375	Aug 01, 2020	DP		
		7342117	Nov 04, 2023	DS		
		7982049	Nov 04, 2023	DP		
		8772315	Oct 30, 2028	U-2300		
		8835474	Nov 04, 2023	U-1527		
		RE44872	Nov 04, 2023	U-1527		
<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>MODAFINIL - PROVIGIL</u>						
N 020717	001	7297346	Nov 29, 2023	DP		
<u>MODAFINIL - PROVIGIL</u>						
N 020717	002	7297346	Nov 29, 2023	DP		
<u>MOMETASONE FUROATE - SINUVA</u>						
N 209310	001	10232152	Nov 24, 2034	DP	U-2272	NP
		10357640	Oct 03, 2031	U-2272		Dec 08, 2020
		10406332	Mar 13, 2034	DP		
		7544192	Nov 29, 2026	U-2272		
		7662141	Mar 12, 2024	U-2272		
		7713255	Mar 12, 2024	U-2272		
		7951130	Mar 12, 2024	U-2272		
		7951131	Mar 12, 2024	U-2272		
		7951133	Mar 12, 2024	U-2272		
		8025635	Jun 12, 2027	DP	U-2272	
		8109918	Mar 12, 2024	U-2272		
		8763222	Feb 08, 2032	DP		
		9585681	Apr 04, 2026	U-2272		
<u>MONTELUKAST SODIUM - SINGULAIR</u>						
N 021409	001	8007830	Oct 24, 2022	DP		
<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		

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<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		
<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		

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<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>MOXIDECTIN - MOXIDECTIN</u>						
N 210867 001					NCE ODE-193	Jun 13, 2023 Jun 13, 2025
<u>MOXIFLOXACIN HYDROCHLORIDE - AVELOX IN SODIUM CHLORIDE 0.8% IN PLASTIC CONTAINER</u>						
N 021277 001	6548079	Jul 25, 2020	DP	U-298		
<u>MOXIFLOXACIN HYDROCHLORIDE - MOXEZA</u>						
N 022428 001	8450311	May 29, 2029	DP			
	9114168	May 29, 2029	DP			
<u>NAFTIFINE HYDROCHLORIDE - NAFTIN</u>						
N 204286 001	10166205	Jan 31, 2033	DP			
	10166206	Jan 31, 2033	DP			
	8778365	Jan 31, 2033	DP			
	9161914	Jan 31, 2033		U-540		
<u>NALDEMEDINE TOSYLATE - SYMPROIC</u>						
N 208854 001	9108975	Nov 11, 2031	DS DP		NCE	Mar 23, 2022
	RE46365	Jan 11, 2028	DS DP			
	RE46375	Oct 05, 2026	DS DP	U-1185		
<u>NALOXEGOL OXALATE - MOVANTIK</u>						
N 204760 001	7056500	Jun 29, 2024	DP	U-1185		
	7662365	Oct 18, 2022	DS DP			
	7786133	Dec 19, 2027	DS DP			
	8067431	Dec 16, 2024		U-1185		
	8617530	Oct 18, 2022		U-1185		
	9012469	Apr 02, 2032	DS DP			
<u>NALOXEGOL OXALATE - MOVANTIK</u>						
N 204760 002	7056500	Jun 29, 2024	DP	U-1185		
	7662365	Oct 18, 2022	DS DP			
	7786133	Dec 19, 2027	DS DP			
	8067431	Dec 16, 2024		U-1185		

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<u>NALOXEGOL OXALATE - MOVANTIK</u>						
N 204760 002	8617530	Oct 18, 2022				U-1185
	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
	10314977	Nov 23, 2024	DP			
	10322239	Feb 28, 2031				U-1907
	10335549	Apr 30, 2025	DP			
	7731686	Jun 10, 2026	DP			
	7731690	Jan 15, 2025	DP			
	7749194	Oct 30, 2028	DP			
	7918823	Nov 23, 2024	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			
	8313466	Nov 23, 2024	DP			
	8361029	Nov 23, 2024	DP			
	8425462	Nov 23, 2024	DP			
	8608698	Nov 23, 2024	DP			
	8627816	Feb 04, 2032	DP			
	8926594	Mar 31, 2026	DP			
	8939943	Feb 28, 2031	DP			
	9022022	Feb 28, 2031	DP			
	9056170	Nov 23, 2024	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
	9737669	Nov 23, 2024	DP			
<u>NALOXONE HYDROCHLORIDE - NARCAN</u>						
N 208411 001	10085937	Mar 16, 2035				U-1903
	9211253	Mar 16, 2035	DP			
	9468747	Mar 16, 2035	DP			U-1903
	9561177	Mar 16, 2035	DP			U-1903
	9629965	Mar 16, 2035	DP			U-1903
	9775838	Mar 16, 2035				U-1903
<u>NALOXONE HYDROCHLORIDE - NARCAN</u>						
N 208411 002	9480644	Mar 16, 2035				DP U-1903
	9707226	Mar 16, 2035				DP U-1903
<u>NALOXONE HYDROCHLORIDE - EVZIO (AUTOINJECTOR)</u>						
N 209862 001	10143792	May 24, 2031				U-2476
	10220158	Mar 20, 2035	DP			U-2500
	10314977	Nov 23, 2024	DP			
	10322239	Feb 28, 2031				U-1907
	10335549	Apr 30, 2025	DP			
	7731686	Jun 01, 2026	DP			
	7731690	Jan 15, 2025	DP			
	7749194	Oct 30, 2028	DP			
	7918823	Nov 23, 2024	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			
	8313466	Nov 23, 2024	DP			
	8361029	Nov 23, 2024	DP			
	8425462	Nov 23, 2024	DP			
	8608698	Nov 23, 2024	DP			
	8627816	Feb 04, 2032	DP			
	8926594	Mar 31, 2026	DP			
	8939943	Feb 28, 2031	DP			

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<u>NALOXONE HYDROCHLORIDE - EVZIO (AUTOINJECTOR)</u>						
N 209862	001	9022022	Feb 28, 2031	DP		
		9056170	Nov 23, 2024	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		
		9737669	Nov 23, 2024	DP		
<u>NALOXONE HYDROCHLORIDE; OXYCODONE HYDROCHLORIDE - TARGINIQ</u>						
N 205777	001	8846090	Apr 04, 2023	DP		
		8846091	Apr 04, 2023	DP		
		8969369	May 10, 2022	DP U-1556		
		9056051	May 10, 2022	DP U-1556		
		9073933	Mar 30, 2025	DS		
		9084729	May 10, 2022	DP U-1556		
		9161937	May 10, 2022	DP U-1556		
		9168252	May 10, 2022	DP U-1556		
		9283216	May 10, 2022	DP U-1819		
		9283221	May 10, 2022	DP U-1819		
		9345701	May 10, 2022	DP U-1819		
		9511066	May 10, 2022	U-1921		
		9522919	Mar 30, 2025	DS DP		
		9555000	Apr 04, 2023	DP U-1556		
		9907793	Apr 04, 2023	DP U-1556		
<u>NALOXONE HYDROCHLORIDE; OXYCODONE HYDROCHLORIDE - TARGINIQ</u>						
N 205777	002	8846090	Apr 04, 2023	DP		
		8846091	Apr 04, 2023	DP		
		8969369	May 10, 2022	DP U-1556		
		9056051	May 10, 2022	DP U-1556		
		9073933	Mar 30, 2025	DS		
		9084729	May 10, 2022	DP U-1556		
		9161937	May 10, 2022	DP U-1556		
		9168252	May 10, 2022	DP U-1556		
		9283216	May 10, 2022	DP U-1819		
		9283221	May 10, 2022	DP U-1819		
		9345701	May 10, 2022	DP U-1819		
		9511066	May 10, 2022	U-1921		
		9522919	Mar 30, 2025	DS DP		
		9555000	Apr 04, 2023	DP U-1556		
		9907793	Apr 04, 2023	DP U-1556		
<u>NALOXONE HYDROCHLORIDE; OXYCODONE HYDROCHLORIDE - TARGINIQ</u>						
N 205777	003	8846090	Apr 04, 2023	DP		
		8846091	Apr 04, 2023	DP		
		8969369	May 10, 2022	DP U-1556		
		9056051	May 10, 2022	DP U-1556		
		9073933	Mar 30, 2025	DS		
		9084729	May 10, 2022	DP U-1556		
		9161937	May 10, 2022	DP U-1556		
		9168252	May 10, 2022	DP U-1556		
		9283216	May 10, 2022	DP U-1819		
		9283221	May 10, 2022	DP U-1819		
		9345701	May 10, 2022	DP U-1819		
		9511066	May 10, 2022	U-1921		
		9522919	Mar 30, 2025	DS DP		
<u>NALTREXONE - VIVITROL</u>						
N 021897	001	6264987	May 19, 2020	DP		
		6379704	May 19, 2020	DP		
		6495164	May 25, 2020	DP		
		6534092	May 19, 2020	DP		
		6667061	May 25, 2020	DP		
		7799345	May 25, 2020	DP		
		7919499	Oct 15, 2029	U-1123		
		7919499	Oct 15, 2029	U-1124		

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<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			
	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 - BYSTOLIC</u>						
N 021742 002	6545040	Dec 17, 2021	DP U-3			
<u>NEBIVOLOL HYDROCHLORIDE - BYSTOLIC</u>						
N 021742 003	6545040	Dec 17, 2021	DP U-3			
<u>NEBIVOLOL HYDROCHLORIDE - BYSTOLIC</u>						
N 021742 004	6545040	Dec 17, 2021	DP U-3			
<u>NEBIVOLOL HYDROCHLORIDE - BYSTOLIC</u>						
N 021742 005	6545040	Dec 17, 2021	DP U-3			
<u>NEBIVOLOL HYDROCHLORIDE; VALSARTAN - BYVALSON</u>						
N 206302 001	7803838	Aug 29, 2026	DP			
	7838552	Oct 04, 2027	U-185			
<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	7947295	Jun 08, 2024	DP			
	8921337	Mar 31, 2032	DP			
	9662398	Dec 01, 2030	DP			
<u>NERATINIB MALEATE - NERLYNX</u>						
N 208051 001	10035788	Oct 15, 2028	U-2043		I-823	Feb 25, 2023
	7399865	Dec 29, 2025	DS DP		NCE	Jul 17, 2022
	7982043	Oct 08, 2025	U-2043			
	8518446	Nov 20, 2030	DP U-2043			
	8790708	Nov 05, 2030	DP U-2043			

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<u>NERATINIB MALEATE - NERLYNX</u>						
N 208051	001	9139558	Oct 15, 2028	U-2043		
		9211291	Mar 24, 2030	U-2043		
		9630946	Oct 15, 2028	U-2043		
<u>NETARSUDIL MESYLATE - RHOPRESSA</u>						
N 208254	001	10174017	Jan 27, 2030	DS DP U-1524	NCE	Dec 18, 2022
		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		
		6297375	Mar 17, 2023	DS		
		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 - CARDENE IN 4.8% DEXTROSE IN PLASTIC CONTAINER</u>						
N 019734	002	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	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 0.83% SODIUM CHLORIDE IN PLASTIC CONTAINER</u>						
N 019734	004	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 - NICODERM CO</u>						
N 020165	004	8075911	May 22, 2021	DP		
		8663680	Feb 13, 2020	DP		
		8999379	Feb 13, 2020		U-1686	
<u>NICOTINE - NICODERM CO</u>						
N 020165	005	8075911	May 22, 2021	DP		
		8663680	Feb 13, 2020	DP		
		8999379	Feb 13, 2020		U-1686	
<u>NICOTINE - NICODERM CO</u>						
N 020165	006	8075911	May 22, 2021	DP		
		8663680	Feb 13, 2020	DP		
		8999379	Feb 13, 2020		U-1686	
<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		

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<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>NILOTINIB HYDROCHLORIDE - TASIGNA</u>						
N 022068 001	7169791	Jul 04, 2023	DS DP U-836		D-170	Dec 22, 2020
	7169791*PED	Jan 04, 2024			NPP	Mar 22, 2021
	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		PED	Jun 22, 2021
	8293756*PED	Mar 25, 2028			PED	Sep 22, 2021
	8389537	Jul 18, 2026	DS DP U-1374		PED	Sep 22, 2025
	8389537*PED	Jan 18, 2027			PED	Sep 22, 2025
	8415363	Jul 18, 2026	DS DP U-1407			
	8415363*PED	Jan 18, 2027				
	8501760	Jul 18, 2026	DS DP			
	8501760*PED	Jan 18, 2027				
	9061029	Apr 07, 2032	DP U-1374			
	9061029*PED	Oct 07, 2032				
<u>NILOTINIB HYDROCHLORIDE - TASIGNA</u>						
N 022068 002	7169791	Jul 04, 2023	DS DP U-836		D-170	Dec 22, 2020
	7169791*PED	Jan 04, 2024			NPP	Mar 22, 2021
	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		PED	Jun 22, 2021
	8293756*PED	Mar 25, 2028			PED	Sep 22, 2021
	8389537	Jul 18, 2026	DS DP U-1374		PED	Sep 22, 2025
	8389537*PED	Jan 18, 2027			PED	Sep 22, 2025
	8415363	Jul 18, 2026	DS DP U-1407			
	8415363*PED	Jan 18, 2027				
	8501760	Jul 18, 2026	DS DP			
	8501760*PED	Jan 18, 2027				
	9061029	Apr 07, 2032	DP U-1374			
	9061029*PED	Oct 07, 2032				
<u>NILOTINIB HYDROCHLORIDE - TASIGNA</u>						
N 022068 003	7169791	Jul 04, 2023	DS DP U-836		NPP	Mar 22, 2021
	7169791*PED	Jan 04, 2024			ODE-171	Mar 22, 2025
	8163904	Aug 23, 2028	DS DP		ODE-172	Mar 22, 2025
	8163904*PED	Feb 23, 2029			PED	Sep 22, 2021
	8293756	Sep 25, 2027	DP		PED	Sep 22, 2025
	8293756*PED	Mar 25, 2028			PED	Sep 22, 2025
	8389537	Jul 18, 2026	DS DP U-1374			
	8389537*PED	Jan 18, 2027				
	8415363	Jul 18, 2026	DS DP U-1407			
	8415363*PED	Jan 18, 2027				
	8501760	Jul 18, 2026	DS DP			
	8501760*PED	Jan 18, 2027				
	9061029	Apr 07, 2032	DS DP U-1374			
	9061029*PED	Oct 07, 2032				
<u>NIMODIPINE - NYMALIZE</u>						
N 203340 001					ODE-46	May 10, 2020
<u>NINTEDANIB ESYLATE - OFEV</u>						
N 205832 001	10105323	Jun 04, 2029	DP		I-805	Sep 06, 2022
	10154990	Dec 20, 2025		U-2620	ODE-261	Sep 06, 2026
	6762180	Oct 03, 2020	DS DP		ODE-77	Oct 15, 2021
	7119093	Feb 21, 2024	DS DP			
	9907756	Jun 07, 2029	DP			
<u>NINTEDANIB ESYLATE - OFEV</u>						
N 205832 002	10105323	Jun 04, 2029	DP		I-805	Sep 06, 2022
	10154990	Dec 20, 2025		U-2620	ODE-261	Sep 06, 2026
	6762180	Oct 03, 2020	DS DP		ODE-77	Oct 15, 2021
	7119093	Feb 21, 2024	DS DP			

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<u>NINTEDANIB ESYLATE - OFEV</u>						
N 205832	002 9907756	Jun 07, 2029	DP			
<u>NIRAPARIB TOSYLATE - ZEJULA</u>						
N 208447	001 8071579	Aug 12, 2027		U-2655	I-813	Oct 23, 2022
	8071623	Mar 22, 2030	DS DP		I-814	Oct 23, 2022
	8143241	Aug 12, 2027		U-2655	NCE	Mar 27, 2022
	8436185	Apr 24, 2029	DS		ODE-133	Mar 27, 2024
	8859562	Aug 04, 2031		U-2655	ODE-277	Oct 23, 2026
					ODE-278	Oct 23, 2026
<u>NITISINONE - ORFADIN</u>						
N 021232	001				D-169	Sep 01, 2020
<u>NITISINONE - ORFADIN</u>						
N 021232	002				D-169	Sep 01, 2020
<u>NITISINONE - ORFADIN</u>						
N 021232	003				D-169	Sep 01, 2020
<u>NITISINONE - ORFADIN</u>						
N 021232	004				D-169	Sep 01, 2020
<u>NITISINONE - ORFADIN</u>						
N 206356	001 9301932	Feb 28, 2033	DP U-1836		D-169	Sep 01, 2020
<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			
<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 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		

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<u>NITRIC OXIDE - INOMAX</u>						
N 020845	003	8846112*PED				
		9265911				
		9265911*PED	Jan 06, 2031	DP U-1824		
		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>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>NIZATIDINE - AXID</u>						
N 021494	001	6930119	Jul 17, 2022	DP		
<u>NUSINERSEN SODIUM - SPINRAZA</u>						
N 209531	001	10266822	Dec 05, 2025	DP U-1942	M-226	May 14, 2021
		10266822	Dec 05, 2025	DP U-1943	NCE	Dec 23, 2021
		10266822	Dec 05, 2025	DP U-1944	ODE-127	Dec 23, 2023
		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		
		7101993	Sep 05, 2023	DS		
		7838657	Jul 11, 2027	DS		
		8110560	Dec 05, 2025	U-1942		
		8110560	Dec 05, 2025	U-1943		
		8110560	Dec 05, 2025	U-1944		
		8361977	May 27, 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	NCE	May 27, 2021
		10052337	Apr 26, 2036	DP	ODE-119	May 27, 2023
		10174073	Jun 17, 2033	DS		
		7138390	Nov 16, 2022	DS DP		
		8058267	Feb 21, 2022	U-1854		
		8377916	Feb 21, 2022	U-1854		
		9238673	Jun 17, 2033	DP		
<u>OBETICHOLIC ACID - OCALIVA</u>						
N 207999	002	10047117	Sep 06, 2033	U-1854	NCE	May 27, 2021
		10052337	Apr 26, 2036	DP	ODE-119	May 27, 2023
		10174073	Jun 17, 2033	DS		
		7138390	Nov 16, 2022	DS DP		
		8058267	Feb 21, 2022	U-1854		
		8377916	Feb 21, 2022	U-1854		
		9238673	Jun 17, 2033	DP		
<u>OCTREOTIDE ACETATE - BYNFEZIA PEN</u>						
N 213224	001	10342850	May 15, 2038	DP		
<u>OLAPARIB - LYNPARZA</u>						
N 206162	001	7151102	Apr 29, 2022	DS DP	ODE-83	Dec 19, 2021
		7449464	Oct 11, 2024	DS DP		
		7981889	Oct 11, 2024	DS DP		
		8143241	Aug 12, 2027	U-1634		

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<u>OLAPARIB - LYNPARZA</u>						
N 206162 001	8247416	Sep 24, 2028	DS			
	8859562	Aug 04, 2031		U-1634		
	8912187	Mar 12, 2024		U-1634		
<u>OLAPARIB - LYNPARZA</u>						
N 208558 001	7151102	Apr 29, 2022	DS DP		I-762	Jan 12, 2021
	7449464	Oct 11, 2024	DS DP		I-776	Dec 19, 2021
	7981889	Oct 11, 2024	DS DP		I-818	Dec 27, 2022
	8143241	Aug 12, 2027		U-2101	NP	Aug 17, 2020
	8143241	Aug 12, 2027		U-2103	ODE-180	Aug 17, 2024
	8143241	Aug 12, 2027		U-2480	ODE-181	Aug 17, 2024
	8143241	Aug 12, 2027		U-2481	ODE-226	Dec 19, 2025
	8143241	Aug 12, 2027		U-2482	ODE-283	Dec 27, 2026
	8143241	Aug 12, 2027		U-2483	ODE-83	Dec 19, 2021
	8143241	Aug 12, 2027		U-2716		
	8475842	Dec 31, 2029	DP			
	8859562	Aug 04, 2031		U-2101		
	8859562	Aug 04, 2031		U-2480		
	8859562	Aug 04, 2031		U-2481		
	8859562	Aug 04, 2031		U-2482		
	8859562	Aug 04, 2031		U-2483		
	8859562	Aug 04, 2031		U-2716		
	8912187	Mar 12, 2024		U-2101		
	8912187	Mar 12, 2024		U-2480		
	8912187	Mar 12, 2024		U-2481		
	8912187	Mar 12, 2024		U-2482		
	8912187	Mar 12, 2024		U-2483		
	9566276	Mar 12, 2024		U-2716		
<u>OLAPARIB - LYNPARZA</u>						
N 208558 002	7151102	Apr 29, 2022	DS DP		I-762	Jan 12, 2021
	7449464	Oct 11, 2024	DS DP		I-776	Dec 19, 2021
	7981889	Oct 11, 2024	DS DP		I-818	Dec 27, 2022
	8143241	Aug 12, 2027		U-2101	NP	Aug 17, 2020
	8143241	Aug 12, 2027		U-2103	ODE-180	Aug 17, 2024
	8143241	Aug 12, 2027		U-2480	ODE-181	Aug 17, 2024
	8143241	Aug 12, 2027		U-2481	ODE-226	Dec 19, 2025
	8143241	Aug 12, 2027		U-2482	ODE-283	Dec 27, 2026
	8143241	Aug 12, 2027		U-2483	ODE-83	Dec 19, 2021
	8143241	Aug 12, 2027		U-2716		
	8475842	Dec 31, 2029	DP			
	8859562	Aug 04, 2031		U-2101		
	8859562	Aug 04, 2031		U-2480		
	8859562	Aug 04, 2031		U-2481		
	8859562	Aug 04, 2031		U-2482		
	8859562	Aug 04, 2031		U-2483		
	8859562	Aug 04, 2031		U-2716		
	8912187	Mar 12, 2024		U-2101		
	8912187	Mar 12, 2024		U-2480		
	8912187	Mar 12, 2024		U-2481		
	8912187	Mar 12, 2024		U-2482		
	8912187	Mar 12, 2024		U-2483		
	9566276	Mar 12, 2024		U-2716		
<u>OLODATEROL HYDROCHLORIDE - STRIVERDI RESPIMAT</u>						
N 203108 001	6988496	Feb 23, 2020		DP U-1547		
	7056916	Dec 07, 2023	DS DP			
	7220742	May 12, 2025	DS DP	U-1547		
	7284474	Aug 26, 2024		DP		
	7396341	Oct 10, 2026		DP U-1547		
	7491719	Nov 10, 2023	DS DP			
	7727984	Jan 19, 2027	DS			
	7786111	Nov 10, 2023		DP		
	7837235	Mar 13, 2028		DP		
	7896264	May 26, 2025		DP		
	7988001	Aug 04, 2021		DP		
	8034809	May 12, 2025		U-1547		
	8044046	Nov 10, 2023		U-1547		
	8733341	Oct 16, 2030		DP		
	9027967	Mar 31, 2027		DP		

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<u>OLODATEROL HYDROCHLORIDE - STRIVERDI RESPIMAT</u>						
N 203108	001	6988496	Feb 23, 2020	DP	U-1547	
		7056916	Dec 07, 2023	DS DP		
		7220742	May 12, 2025	DS DP	U-1547	
		7284474	Aug 26, 2024	DP		
		7396341	Oct 10, 2026	DP	U-1547	
		7491719	Nov 10, 2023	DS DP		
		7727984	Jan 19, 2027	DS		
		7786111	Nov 10, 2023	DP		
		7837235	Mar 13, 2028	DP		
		7896264	May 26, 2025	DP		
		7988001	Aug 04, 2021	DP		
		8034809	May 12, 2025		U-1547	
		8044046	Nov 10, 2023		U-1547	
		8733341	Oct 16, 2030	DP		
		9027967	Mar 31, 2027	DP		
<u>OLODATEROL HYDROCHLORIDE; TIOTROPIUM BROMIDE - STIOLTO RESPIMAT</u>						
N 206756	001	6988496	Feb 23, 2020	DP		M-233 Oct 05, 2021
		6988496*PED	Aug 23, 2020			
		7056916	Dec 07, 2023	DS DP		
		7220742	May 12, 2025	DS DP	U-1703	
		7284474	Aug 26, 2024	DP		
		7284474*PED	Feb 26, 2025			
		7396341	Oct 10, 2026	DP		
		7396341*PED	Apr 10, 2027			
		7491719	Nov 10, 2023	DS DP		
		7727984	Jan 19, 2027	DS		
		7786111	Nov 10, 2023	DP		
		7837235	Mar 13, 2028	DP		
		7837235*PED	Sep 13, 2028			
		7896264	May 26, 2025	DP		
		7988001	Aug 04, 2021	DP		
		8034809	May 12, 2025		U-1702	
		8044046	Nov 10, 2023		U-1702	
		8733341	Oct 16, 2030	DP		
		9027967	Mar 31, 2027	DP		
<u>OLOPATADINE HYDROCHLORIDE - PATADAY</u>						
N 021545	001	6995186	Nov 12, 2023	DP	U-765	
		7402609	Jun 19, 2022	DP		
<u>OLOPATADINE HYDROCHLORIDE - PATANASE</u>						
N 021861	001	7977376	Feb 02, 2023	DP		
		8399508	Sep 17, 2022		U-726	
		8399508*PED	Mar 17, 2023			
<u>OLOPATADINE HYDROCHLORIDE - PAZEO</u>						
N 206276	001	8791154	May 19, 2032	DP	U-1680	
		9533053	May 19, 2032	DP		
<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	
		7326696	Sep 24, 2023	DS		
		7553828	Jun 02, 2023	DS		
		8383610	Sep 23, 2030	DS		
		9265740	Mar 05, 2029		U-1569	
		9314475	Mar 18, 2031	DP		
		9365500	Jun 29, 2021		U-1569	
		9365500	Jun 29, 2021		U-2576	
		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
		7326696	Sep 24, 2023	DS		

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<u>OMADACYCLINE TOSYLATE - NUZYRA</u>						
N 209817	001 7553828	Jun 02, 2023	DS			
	9265740	Mar 05, 2029	DP			
	9365500	Jun 29, 2021		U-1569		
	9365500	Jun 29, 2021		U-2576		
	9724358	Mar 05, 2029		U-1569		
<u>OMBITASVIR; PARITAPREVIR; RITONAVIR - TECHNIVIE</u>						
N 207931	001 7148359*PED	Jan 19, 2020				
	7364752	Nov 10, 2020	DP			
	7364752*PED	May 10, 2021				
	8268349	Aug 25, 2024	DP			
	8268349*PED	Feb 25, 2025				
	8399015	Aug 25, 2024	DP			
	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	
	5792795	May 13, 2020	DP			
	5948818	May 13, 2020	DP			
	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			
<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>ORITAVANCIN DIPHOSPHATE - ORBACTIV</u>						
N 206334	001 5840684	Nov 24, 2020	DS DP	U-1569	NCE	Aug 06, 2019
	8420592	Aug 29, 2029		U-1570	GAIN	Aug 06, 2024
	9649352	Jul 16, 2035	DP			
	9682061	Apr 26, 2030		U-1569		
<u>OSELTAMIVIR PHOSPHATE - TAMIFLU</u>						
N 021087	001				M-251	Aug 02, 2022
<u>OSELTAMIVIR PHOSPHATE - TAMIFLU</u>						
N 021087	002				M-251	Aug 02, 2022
<u>OSELTAMIVIR PHOSPHATE - TAMIFLU</u>						
N 021087	003				M-251	Aug 02, 2022
<u>OSELTAMIVIR PHOSPHATE - TAMIFLU</u>						
N 021246	001				M-251	Aug 02, 2022
<u>OSELTAMIVIR PHOSPHATE - TAMIFLU</u>						
N 021246	002				M-251	Aug 02, 2022

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<u>OSIMERTINIB MESYLATE - TAGRISSO</u>						
N 208065 001	10183020	Jan 02, 2035	DP U-1777		I-774	Apr 18, 2021
	10183020	Jan 02, 2035	DP U-2289		NCE	Nov 13, 2020
	8946235	Aug 08, 2032	DS DP U-1777		ODE-102	Nov 13, 2022
	8946235	Aug 08, 2032	DS DP U-2289		ODE-176	Apr 18, 2025
	9732058	Jul 25, 2032	DS DP U-1777			
	9732058	Jul 25, 2032	DS DP U-2289			
<u>OSIMERTINIB MESYLATE - TAGRISSO</u>						
N 208065 002	10183020	Jan 02, 2035	DP U-1777		I-774	Apr 18, 2021
	10183020	Jan 02, 2035	DP U-2289		NCE	Nov 13, 2020
	8946235	Aug 08, 2032	DS DP U-1777		ODE-102	Nov 13, 2022
	8946235	Aug 08, 2032	DS DP U-2289		ODE-176	Apr 18, 2025
	9732058	Jul 25, 2032	DS DP U-1777			
	9732058	Jul 25, 2032	DS DP U-2289			
<u>OSPEMIFENE - OSPHENA</u>						
N 203505 001	6245819	Jul 21, 2025	U-1370		I-793	Jan 25, 2022
	6245819	Jul 21, 2025	U-905			
	8236861	Aug 11, 2026	U-1369			
	8236861	Aug 11, 2026	U-1370			
	8236861	Aug 11, 2026	U-905			
	8470890	Feb 13, 2024	U-1369			
	8470890	Feb 13, 2024	U-1370			
	8470890	Feb 13, 2024	U-905			
	8642079	Jul 09, 2028	DP			
	8772353	Feb 13, 2024	U-1369			
	8772353	Feb 13, 2024	U-1370			
	8772353	Feb 13, 2024	U-905			
	9241915	Feb 13, 2024	U-1369			
	9241915	Feb 13, 2024	U-1370			
	9241915	Feb 13, 2024	U-905			
	9566252	Jul 21, 2020	U-1370			
	9855224	Feb 13, 2024	U-1369			
	9855224	Feb 13, 2024	U-1370			
	9855224	Feb 13, 2024	U-905			
<u>OXCARBAZEPINE - TRILEPTAL</u>						
N 021285 001	8119148	Dec 19, 2020	DP U-724			
<u>OXCARBAZEPINE - OXTELLAR XR</u>						
N 202810 001	10220042	Apr 13, 2027	U-2501			
	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			
	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			
	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			

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<u>OXCARBAZEPINE - OXTELLAR XR</u>						
N 202810	003 9855278	Apr 13, 2027	DP			
<u>OXYBUTYNYNIN - OXYTROL</u>						
N 021351	002 6743441	Apr 26, 2020	DP U-318			
	7081249	Apr 26, 2020	DP U-318			
	7081250	Apr 26, 2020	DP U-318			
	7081251	Apr 26, 2020	DP U-318			
	7081252	Apr 26, 2020	DP U-318			
	7179483	Apr 26, 2020	DS DP U-318			
<u>OXYBUTYNYNIN - OXYTROL FOR WOMEN</u>						
N 202211	001 6743441	Apr 26, 2020	DP U-1329			
	7081249	Apr 26, 2020	DP U-1329			
	7081250	Apr 26, 2020	DP U-1329			
	7081251	Apr 26, 2020	DP U-1329			
	7081252	Apr 26, 2020	DP U-1329			
	7179483	Apr 26, 2020	U-1329			
<u>OXYBUTYNYNIN - GELNIQUE 3%</u>						
N 202513	001 7029694	Apr 26, 2020	DP U-318			
	7179483	Apr 26, 2020	U-318			
	7198801	Jun 25, 2022	DP			
	8241662	Apr 26, 2020	U-318			
<u>OXYBUTYNYNIN CHLORIDE - GELNIQUE</u>						
N 022204	001 10272061	Apr 26, 2020		U-2525		
	10449173	Nov 06, 2029	DP U-2637			
	7029694	Apr 26, 2020	DP U-318			
	7179483	Apr 26, 2020	U-318			
	8241662	Apr 26, 2020	U-318			
	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			
	10525052	Jul 07, 2023	DP U-1556			
	10525053	Jul 07, 2023	DP			
	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			
	8840928	Jul 07, 2023	DP U-1556			
	9044398	Jul 07, 2023	DP			
	9248195	Jul 07, 2023	U-1556			
	9592200	Jul 07, 2023	DP			
	9682075	Dec 10, 2030	DP U-1556			
	9737530	Sep 02, 2036	DP U-1556			
	9763883	Jul 07, 2023	DP			
	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			
	10525052	Jul 07, 2023	DP U-1556			
	10525053	Jul 07, 2023	DP			
	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			
	8840928	Jul 07, 2023	DP U-1556			
	9044398	Jul 07, 2023	DP			
	9248195	Jul 07, 2023	U-1556			
	9592200	Jul 07, 2023	DP			
	9682075	Dec 10, 2030	DP U-1556			
	9737530	Sep 02, 2036	DP U-1556			
	9763883	Jul 07, 2023	DP			
	9968598	Sep 02, 2036	DP U-1556			

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<u>OXYCODONE - XTAMPZA ER</u>						
N 208090 002	10004729	Dec 10, 2030	DP U-1556			
	10188644	Sep 02, 2036	DP U-1556			
	10525052	Jul 07, 2023	DP U-1556			
	10525053	Jul 07, 2023	DP			
	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			
	8840928	Jul 07, 2023	DP U-1556			
	9044398	Jul 07, 2023	DP			
	9248195	Jul 07, 2023	U-1556			
	9592200	Jul 07, 2023	DP			
	9682075	Dec 10, 2030	DP U-1556			
	9737530	Sep 02, 2036	DP U-1556			
	9763883	Jul 07, 2023	DP			
	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			
	10525052	Jul 07, 2023	DP U-1556			
	10525053	Jul 07, 2023	DP			
	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			
	8840928	Jul 07, 2023	DP U-1556			
	9044398	Jul 07, 2023	DP			
	9248195	Jul 07, 2023	U-1556			
	9592200	Jul 07, 2023	DP			
	9682075	Dec 10, 2030	DP U-1556			
	9737530	Sep 02, 2036	DP U-1556			
	9763883	Jul 07, 2023	DP			
	9968598	Sep 02, 2036	DP U-1556			
<u>OXYCODONE - XTAMPZA ER</u>						
N 208090 004	10004729	Dec 10, 2030	DP U-1556			
	10188644	Sep 02, 2036	DP U-1556			
	10525052	Jul 07, 2023	DP U-1556			
	10525053	Jul 07, 2023	DP			
	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			
	8840928	Jul 07, 2023	DP U-1556			
	9044398	Jul 07, 2023	DP			
	9248195	Jul 07, 2023	U-1556			
	9592200	Jul 07, 2023	DP			
	9682075	Dec 10, 2030	DP U-1556			
	9737530	Sep 02, 2036	DP U-1556			
	9763883	Jul 07, 2023	DP			
	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			
	10525052	Jul 07, 2023	DP U-1556			
	10525053	Jul 07, 2023	DP			
	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			
	8840928	Jul 07, 2023	DP U-1556			
	9044398	Jul 07, 2023	DP			
	9248195	Jul 07, 2023	U-1556			
	9592200	Jul 07, 2023	DP			

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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 005	9682075	Dec 10, 2030	DP U-1556			
	9737530	Sep 02, 2036	DP U-1556			
	9763883	Jul 07, 2023	DP			
	9968598	Sep 02, 2036	DP U-1556			
<u>OXYCODONE HYDROCHLORIDE - OXYCONTIN</u>						
N 022272 001	10130591	Nov 20, 2023	DP U-1819			
	10369109	Jun 16, 2023	DP			
	10407434	Mar 30, 2025	DS			
	7674799	Mar 30, 2025	DP		Y	
	7674800	Mar 30, 2025	DS		Y	
	7683072	Mar 30, 2025	DS		Y	
	7776314	Apr 19, 2025	DP		Y	
	8309060	Nov 20, 2023	DP U-1556			
	8808741	Aug 24, 2027	U-1556			
	8894987	Mar 29, 2030	DP			
	8894988	Aug 24, 2027	DP			
	9060976	Aug 06, 2022	DP		Y	
	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			
	9675610	Jun 16, 2023	DP			
	9763933	Aug 24, 2027	DP			
	9770416	Aug 24, 2027	DP			
	9775808	Aug 24, 2027	DP			
<u>OXYCODONE HYDROCHLORIDE - OXYCONTIN</u>						
N 022272 002	10130591	Nov 20, 2023	DP U-1819			
	10369109	Jun 16, 2023	DP			
	10407434	Mar 30, 2025	DS			
	7674799	Mar 30, 2025	DP		Y	
	7674800	Mar 30, 2025	DS		Y	
	7683072	Mar 30, 2025	DS		Y	
	7776314	Apr 19, 2025	DP		Y	
	8309060	Nov 20, 2023	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			
	9675610	Jun 16, 2023	DP			
	9763933	Aug 24, 2027	DP			
	9770416	Aug 24, 2027	DP			
	9775808	Aug 24, 2027	DP			
<u>OXYCODONE HYDROCHLORIDE - OXYCONTIN</u>						
N 022272 003	10130591	Nov 20, 2023	DP U-1819			
	10369109	Jun 16, 2023	DP			
	10407434	Mar 30, 2025	DS			
	7674799	Mar 30, 2025	DP		Y	
	7674800	Mar 30, 2025	DS		Y	
	7683072	Mar 30, 2025	DS		Y	
	7776314	Apr 19, 2025	DP		Y	
	8309060	Nov 20, 2023	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			

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<u>OXYCODONE HYDROCHLORIDE - OXYCONTIN</u>						
N 022272	003	9675610	Jun 16, 2023	DP		
		9763933	Aug 24, 2027	DP		
		9770416	Aug 24, 2027	DP		
		9775808	Aug 24, 2027	DP		
<u>OXYCODONE HYDROCHLORIDE - OXYCONTIN</u>						
N 022272	004	10130591	Nov 20, 2023	DP	U-1819	
		10369109	Jun 16, 2023	DP		
		10407434	Mar 30, 2025	DS		
		8309060	Nov 20, 2023	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	
		9675610	Jun 16, 2023	DP		
		9763933	Aug 24, 2027	DP		
		9770416	Aug 24, 2027	DP		
		9775808	Aug 24, 2027	DP		
<u>OXYCODONE HYDROCHLORIDE - OXYCONTIN</u>						
N 022272	005	10130591	Nov 20, 2023	DP	U-1819	
		10369109	Jun 16, 2023	DP		
		10407434	Mar 30, 2025	DS		
		8309060	Nov 20, 2023	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	
		9675610	Jun 16, 2023	DP		
		9763933	Aug 24, 2027	DP		
		9770416	Aug 24, 2027	DP		
		9775808	Aug 24, 2027	DP		
<u>OXYCODONE HYDROCHLORIDE - OXYCONTIN</u>						
N 022272	006	10130591	Nov 20, 2023	DP	U-1819	
		10369109	Jun 16, 2023	DP		
		10407434	Mar 30, 2025	DS		
		8309060	Nov 20, 2023	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	
		9675610	Jun 16, 2023	DP		
		9763933	Aug 24, 2027	DP		
		9770416	Aug 24, 2027	DP		
		9775808	Aug 24, 2027	DP		
<u>OXYCODONE HYDROCHLORIDE - OXYCONTIN</u>						
N 022272	007	10130591	Nov 20, 2023	DP	U-1819	
		10369109	Jun 16, 2023	DP		
		10407434	Mar 30, 2025	DS		
		8309060	Nov 20, 2023	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	

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<u>OXYCODONE HYDROCHLORIDE - OXYCONTIN</u>						
N 022272	007	9492392	Aug 24, 2027	DP		
		9492393	Aug 24, 2027		U-1556	
		9522919	Mar 30, 2025	DS DP		
		9675610	Jun 16, 2023	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		
		7510726	Nov 26, 2023	DP		
		7981439	Nov 26, 2023	DP		
		8409616	Nov 26, 2023	DP		
		8637540	Nov 26, 2023	DP		
		9492443	May 26, 2024	DP		
<u>OXYCODONE HYDROCHLORIDE - OXAYDO</u>						
N 202080	002	7201920	Mar 16, 2025	DP		
		7510726	Nov 26, 2023	DP		
		7981439	Nov 26, 2023	DP		
		8409616	Nov 26, 2023	DP		
		8637540	Nov 26, 2023	DP		
		9492443	May 26, 2024	DP		
<u>OXYCODONE HYDROCHLORIDE - ROXYBOND</u>						
N 209777	001	10314788	Aug 12, 2028	DP	NP	Apr 20, 2020
		7955619	Aug 12, 2028	DP		
<u>OXYCODONE HYDROCHLORIDE - ROXYBOND</u>						
N 209777	002	10314788	Aug 12, 2028	DP	NP	Apr 20, 2020
		7955619	Aug 12, 2028	DP		
<u>OXYCODONE HYDROCHLORIDE - ROXYBOND</u>						
N 209777	003	10314788	Aug 12, 2028	DP	NP	Apr 20, 2020
		7955619	Aug 12, 2028	DP		
<u>OXYMETAZOLINE HYDROCHLORIDE - RHOFADÉ</u>						
N 208552	001	10335391	Jun 11, 2035		U-2567	NP
		7812049	May 02, 2028		U-1959	Jan 18, 2020
		8420688	Aug 02, 2024		U-1959	
		8815929	Jan 22, 2024		U-1959	
		8883838	Dec 01, 2031	DP		
		9974773	Jun 11, 2035		U-2306	
<u>OXYMETAZOLINE HYDROCHLORIDE; TETRACAINE HYDROCHLORIDE - KOVANAZE</u>						
N 208032	001	6413499	Mar 20, 2020		U-1876	
		8580282	Apr 02, 2030	DP	U-1876	
		9308191	Apr 02, 2030	DP	U-1876	
<u>OXYMORPHONE HYDROCHLORIDE - OPANA ER</u>						
N 021610	001	7276250	Feb 04, 2023	DP	U-826	
<u>OXYMORPHONE HYDROCHLORIDE - OPANA ER</u>						
N 021610	002	7276250	Feb 04, 2023	DP	U-826	
<u>OXYMORPHONE HYDROCHLORIDE - OPANA ER</u>						
N 021610	003	7276250	Feb 04, 2023	DP	U-826	
<u>OXYMORPHONE HYDROCHLORIDE - OPANA ER</u>						
N 021610	004	7276250	Feb 04, 2023	DP	U-826	
<u>OXYMORPHONE HYDROCHLORIDE - OPANA ER</u>						
N 021610	005	7276250	Feb 04, 2023	DP	U-826	
<u>OXYMORPHONE HYDROCHLORIDE - OPANA ER</u>						
N 021610	006	7276250	Feb 04, 2023	DP	U-826	
<u>OXYMORPHONE HYDROCHLORIDE - OPANA ER</u>						
N 021610	007	7276250	Feb 04, 2023	DP	U-826	

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<u>OXYMORPHONE HYDROCHLORIDE - OPANA ER</u>						
N 201655 001	7851482	Jul 10, 2029	DS			
	8075872	Nov 20, 2023	DP			
	8114383	Aug 08, 2024	DP			
	8192722	Sep 15, 2025	DP			
	8309060	Nov 20, 2023	DP			
	8309122	Feb 04, 2023	DP			
	8329216	Feb 04, 2023	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			
	8075872	Nov 20, 2023	DP			
	8114383	Aug 08, 2024	DP			
	8192722	Sep 15, 2025	DP			
	8309060	Nov 20, 2023	DP			
	8309122	Feb 04, 2023	DP			
	8329216	Feb 04, 2023	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			
	8075872	Nov 20, 2023	DP			
	8114383	Aug 08, 2024	DP			
	8192722	Sep 15, 2025	DP			
	8309060	Nov 20, 2023	DP			
	8309122	Feb 04, 2023	DP			
	8329216	Feb 04, 2023	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			
	8075872	Nov 20, 2023	DP			
	8114383	Aug 08, 2024	DP			
	8192722	Sep 15, 2025	DP			
	8309060	Nov 20, 2023	DP			
	8309122	Feb 04, 2023	DP			
	8329216	Feb 04, 2023	DP			
	8808737	Jun 21, 2027		U-1598		
	8871779	Nov 22, 2029	DS			
<u>OXYMORPHONE HYDROCHLORIDE - OPANA ER</u>						
N 201655 005	7851482	Jul 10, 2029	DS			
	8075872	Nov 20, 2023	DP			
	8114383	Aug 08, 2024	DP			
	8192722	Sep 15, 2025	DP			
	8309060	Nov 20, 2023	DP			
	8309122	Feb 04, 2023	DP			
	8329216	Feb 04, 2023	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			
	8075872	Nov 20, 2023	DP			
	8114383	Aug 08, 2024	DP			
	8192722	Sep 15, 2025	DP			
	8309060	Nov 20, 2023	DP			
	8309122	Feb 04, 2023	DP			
	8329216	Feb 04, 2023	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			
	8075872	Nov 20, 2023	DP			
	8114383	Aug 08, 2024	DP			
	8192722	Sep 15, 2025	DP			

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<u>OXYMORPHONE HYDROCHLORIDE - OPANA ER</u>						
N 201655 007	8309060	Nov 20, 2023	DP			
	8309122	Feb 04, 2023	DP			
	8329216	Feb 04, 2023	DP			
	8808737	Jun 21, 2027		U-1598		
	8871779	Nov 22, 2029	DS			
<u>OZENOXACIN - XEPI</u>						
N 208945 001	6335447	Apr 06, 2021	DS		NCE	Dec 11, 2022
	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	ODE-52	Sep 06, 2020
	7758891*PED	Aug 21, 2026			PED	Mar 06, 2021
	7820788	Oct 27, 2024	DP	U-1092		
	7820788	Oct 27, 2024	DP	U-1290		
	7820788	Oct 27, 2024	DP	U-1434		
	7820788*PED	Apr 27, 2025				
	7923536	Dec 09, 2023		U-1117		
	7923536	Dec 09, 2023		U-1290		
	7923536	Dec 09, 2023		U-1434		
	7923536*PED	Jun 09, 2024				
	8034375	Aug 13, 2026		U-1290		
	8138229	Dec 09, 2023	DP	U-1092		
	8138229	Dec 09, 2023	DP	U-1290		
	8138229	Dec 09, 2023	DP	U-1434		
	8138229*PED	Jun 09, 2024				
	8268348	Feb 21, 2026		U-1290		
	8314156	Dec 09, 2023		U-1290		
	8314156	Dec 09, 2023		U-1434		
	8314156*PED	Jun 09, 2024				
	8853260	Oct 10, 2020	DP	U-1092		
	8853260	Oct 10, 2020	DP	U-1290		
	8853260	Oct 10, 2020	DP	U-1434		
	8853260*PED	Apr 10, 2021				
	9101543	Feb 21, 2026		U-1434		
	9101543*PED	Aug 21, 2026				
	9393318	Mar 04, 2032		U-1290		
	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>PALBOCICLIB - IBRANCE</u>						
N 207103 001	6936612	Jan 22, 2023	DS DP		NCE	Feb 03, 2020
	7208489	Jan 16, 2023	DS DP			
	7456168	Jan 16, 2023		U-1998		
	7456168	Jan 16, 2023		U-2515		
	RE47739	Jan 16, 2023	DS DP			
<u>PALBOCICLIB - IBRANCE</u>						
N 207103 002	6936612	Jan 22, 2023	DS DP		NCE	Feb 03, 2020
	7208489	Jan 16, 2023	DS DP			
	7456168	Jan 16, 2023		U-1998		
	7456168	Jan 16, 2023		U-2515		
	RE47739	Jan 16, 2023	DS DP			
<u>PALBOCICLIB - IBRANCE</u>						
N 207103 003	6936612	Jan 22, 2023	DS DP		NCE	Feb 03, 2020
	7208489	Jan 16, 2023	DS DP			
	7456168	Jan 16, 2023		U-1998		
	7456168	Jan 16, 2023		U-2515		
	RE47739	Jan 16, 2023	DS DP			
<u>PALBOCICLIB - IBRANCE</u>						
N 212436 001	6936612	Jan 22, 2023	DS DP		NCE	Feb 03, 2020
	7456168	Jan 16, 2023		U-2515		
	RE47739	Jan 16, 2023	DS DP			

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<u>PALBOCICLIB - IBRANCE</u>						
N 212436 002	6936612	Jan 22, 2023	DS DP		NCE	Feb 03, 2020
	7456168	Jan 16, 2023		U-2515		
	RE47739	Jan 16, 2023	DS DP			
<u>PALBOCICLIB - IBRANCE</u>						
N 212436 003	6936612	Jan 22, 2023	DS DP		NCE	Feb 03, 2020
	7456168	Jan 16, 2023		U-2515		
	RE47739	Jan 16, 2023	DS DP			
<u>PALIPERIDONE PALMITATE - INVEGA SUSTENNA</u>						
N 022264 001	9439906	Jan 26, 2031		U-1901	M-215	Dec 20, 2020
	9439906	Jan 26, 2031		U-543		
<u>PALIPERIDONE PALMITATE - INVEGA SUSTENNA</u>						
N 022264 002	9439906	Jan 26, 2031		U-1901	M-215	Dec 20, 2020
	9439906	Jan 26, 2031		U-543		
<u>PALIPERIDONE PALMITATE - INVEGA SUSTENNA</u>						
N 022264 003	9439906	Jan 26, 2031		U-1901	M-215	Dec 20, 2020
	9439906	Jan 26, 2031		U-543		
<u>PALIPERIDONE PALMITATE - INVEGA SUSTENNA</u>						
N 022264 004	9439906	Jan 26, 2031		U-1901	M-215	Dec 20, 2020
	9439906	Jan 26, 2031		U-543		
<u>PALIPERIDONE PALMITATE - INVEGA SUSTENNA</u>						
N 022264 005	9439906	Jan 26, 2031		U-1901	M-215	Dec 20, 2020
	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>PALONOSETRON HYDROCHLORIDE - ALOXI</u>						
N 021372 001	7947724	Jan 30, 2024	DP			
	7947724*PED	Jul 30, 2024				
	7947725	Jan 30, 2024	DP			
	7947725*PED	Jul 30, 2024				
	7960424	Jan 30, 2024	DP			
	7960424*PED	Jul 30, 2024				
	8518981	Jan 30, 2024	DP			
	8518981*PED	Jul 30, 2024				
	8598218	Jan 30, 2024	DP			
	8598218*PED	Jul 30, 2024				
	8598219	Jan 30, 2024	DP			
	8598219*PED	Jul 30, 2024				
	8729094	Jan 30, 2024	DP U-528			
	8729094*PED	Jul 30, 2024				
	9066980	Jan 30, 2024	DP U-528			
	9066980*PED	Jul 30, 2024				
	9125905	Jan 30, 2024	DP			
	9125905*PED	Jul 30, 2024				
	9173942	Jan 30, 2024	DP			
	9173942*PED	Jul 30, 2024				
	9439854	Jan 30, 2024	DP			
	9439854*PED	Jul 30, 2024				
	9457020	Jan 30, 2024	DP			
	9457020*PED	Jul 30, 2024				

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APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>PALONOSETRON HYDROCHLORIDE - ALOXI</u>						
N 021372	001	9457021	Jan 30, 2024	DP		
		9457021*PED	Jul 30, 2024			
<u>PALONOSETRON HYDROCHLORIDE - ALOXI</u>						
N 021372	002	7947724	Jan 30, 2024	DP		
		7947724*PED	Jul 30, 2024			
		7947725	Jan 30, 2024	DP		
		7947725*PED	Jul 30, 2024			
		7960424	Jan 30, 2024	DP		
		7960424*PED	Jul 30, 2024			
		8518981	Jan 30, 2024	DP		
		8518981*PED	Jul 30, 2024			
		8598218	Jan 30, 2024	DP		
		8598218*PED	Jul 30, 2024			
		9173942	Jan 30, 2024	DP		
		9173942*PED	Jul 30, 2024			
		9439854	Jan 30, 2024	DP		
		9439854*PED	Jul 30, 2024			
		9457020	Jan 30, 2024	DP		
		9457020*PED	Jul 30, 2024			
<u>PANCRELIPASE (AMYLASE;LIPASE;PROTEASE) - ZENPEP</u>						
N 022210	001	7658918	Feb 20, 2028	DP		
		8221747	Feb 20, 2028	DP		
		8246950	Feb 20, 2028		U-1274	
		8562978	Feb 20, 2028	DP		
		8562979	Feb 20, 2028	DP	U-1274	
		8562980	Feb 20, 2028	DP	U-1274	
		8562981	Feb 20, 2028		U-1274	
<u>PANCRELIPASE (AMYLASE;LIPASE;PROTEASE) - ZENPEP</u>						
N 022210	002	7658918	Feb 20, 2028	DP		
		8221747	Feb 20, 2028	DP		
		8246950	Feb 20, 2028		U-1274	
		8562978	Feb 20, 2028	DP		
		8562979	Feb 20, 2028	DP	U-1274	
		8562980	Feb 20, 2028	DP	U-1274	
		8562981	Feb 20, 2028		U-1274	
<u>PANCRELIPASE (AMYLASE;LIPASE;PROTEASE) - ZENPEP</u>						
N 022210	003	7658918	Feb 20, 2028	DP		
		8221747	Feb 20, 2028	DP		
		8246950	Feb 20, 2028		U-1274	
		8562978	Feb 20, 2028	DP		
		8562979	Feb 20, 2028	DP	U-1274	
		8562980	Feb 20, 2028	DP	U-1274	
		8562981	Feb 20, 2028		U-1274	
<u>PANCRELIPASE (AMYLASE;LIPASE;PROTEASE) - ZENPEP</u>						
N 022210	004	7658918	Feb 20, 2028	DP		
		8221747	Feb 20, 2028	DP		
		8246950	Feb 20, 2028		U-1274	
		8562978	Feb 20, 2028	DP		
		8562979	Feb 20, 2028	DP	U-1274	
		8562980	Feb 20, 2028	DP	U-1274	
		8562981	Feb 20, 2028		U-1274	
<u>PANCRELIPASE (AMYLASE;LIPASE;PROTEASE) - ZENPEP</u>						
N 022210	005	8221747	Feb 20, 2028	DP		
		8562978	Feb 20, 2028	DP		
		8562979	Feb 20, 2028	DP	U-1274	
		8562980	Feb 20, 2028	DP	U-1274	
		8562981	Feb 20, 2028		U-1274	
<u>PANCRELIPASE (AMYLASE;LIPASE;PROTEASE) - ZENPEP</u>						
N 022210	006	8221747	Feb 20, 2028	DP		
		8562978	Feb 20, 2028	DP		
		8562979	Feb 20, 2028	DP	U-1274	
		8562980	Feb 20, 2028	DP	U-1274	
		8562981	Feb 20, 2028		U-1274	

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<u>PANCRELIPASE (AMYLASE;LIPASE;PROTEASE) - ZENPEP</u>						
N 022210 006	8221747	Feb 20, 2028	DP			
	8562978	Feb 20, 2028	DP			
	8562979	Feb 20, 2028	DP	U-1274		
	8562980	Feb 20, 2028	DP	U-1274		
	8562981	Feb 20, 2028		U-1274		
<u>PANCRELIPASE (AMYLASE;LIPASE;PROTEASE) - ZENPEP</u>						
N 022210 007	8221747	Feb 20, 2028	DP			
	8562978	Feb 20, 2028	DP			
	8562979	Feb 20, 2028	DP	U-1274		
	8562980	Feb 20, 2028	DP	U-1274		
	8562981	Feb 20, 2028		U-1274		
<u>PANCRELIPASE (AMYLASE;LIPASE;PROTEASE) - PANCREAZE</u>						
N 022523 005	8221747	Feb 20, 2028	DP			
	8562978	Feb 20, 2028	DP			
	8562979	Feb 20, 2028	DP	U-1274		
	8562980	Feb 20, 2028	DP	U-1274		
	8562981	Feb 20, 2028	DP	U-1274		
<u>PANCRELIPASE (AMYLASE;LIPASE;PROTEASE); PANCRELIPASE (AMYLASE;LIPASE;PROTEASE); PANCRELIPASE (AMYLASE;LIPASE;PROTEASE) - CREON</u>						
N 020725 001	9198871	Feb 07, 2030	DP	U-1787		
<u>PANCRELIPASE (AMYLASE;LIPASE;PROTEASE); PANCRELIPASE (AMYLASE;LIPASE;PROTEASE); PANCRELIPASE (AMYLASE;LIPASE;PROTEASE) - CREON</u>						
N 020725 002	9198871	Feb 07, 2030	DP	U-1787		
<u>PANCRELIPASE (AMYLASE;LIPASE;PROTEASE); PANCRELIPASE (AMYLASE;LIPASE;PROTEASE); PANCRELIPASE (AMYLASE;LIPASE;PROTEASE) - CREON</u>						
N 020725 003	9198871	Feb 07, 2030	DP	U-1787		
<u>PANCRELIPASE (AMYLASE;LIPASE;PROTEASE); PANCRELIPASE (AMYLASE;LIPASE;PROTEASE); PANCRELIPASE (AMYLASE;LIPASE;PROTEASE) - CREON</u>						
N 020725 004	9198871	Feb 07, 2030	DP	U-1787		
<u>PANCRELIPASE (AMYLASE;LIPASE;PROTEASE); PANCRELIPASE (AMYLASE;LIPASE;PROTEASE); PANCRELIPASE (AMYLASE;LIPASE;PROTEASE) - CREON</u>						
N 020725 005	9198871	Feb 07, 2030	DP	U-1787		
<u>PANOBINOSTAT LACTATE - FARYDAK</u>						
N 205353 001	6552065	Aug 31, 2021	DS DP		NCE	Feb 23, 2020
	6833384	Sep 30, 2021	DS DP	U-1669	ODE-89	Feb 23, 2022
	7067551	Aug 31, 2021		U-1669		
	7989494	Jan 17, 2028	DS DP			
	8883842	Jun 13, 2028		U-1669		
<u>PANOBINOSTAT LACTATE - FARYDAK</u>						
N 205353 002	6552065	Aug 31, 2021	DS DP		NCE	Feb 23, 2020
	6833384	Sep 30, 2021	DS DP	U-1669	ODE-89	Feb 23, 2022
	7067551	Aug 31, 2021		U-1669		
	7989494	Jan 17, 2028	DS DP			
	8883842	Jun 13, 2028		U-1669		
<u>PANOBINOSTAT LACTATE - FARYDAK</u>						
N 205353 003	6552065	Aug 31, 2021	DS DP		NCE	Feb 23, 2020
	6833384	Sep 30, 2021	DS DP	U-1669	ODE-89	Feb 23, 2022
	7067551	Aug 31, 2021		U-1669		
	7989494	Jan 17, 2028	DS DP			
	8883842	Jun 13, 2028		U-1669		
<u>PANTOPRAZOLE SODIUM - PROTONIX IV</u>						
N 020988 001	6780881	Nov 17, 2021	DP			
	7351723	Nov 17, 2021	DP			
	8754108	Nov 17, 2021	DP			
	8754108*PED	May 17, 2022				
<u>PANTOPRAZOLE SODIUM - PROTONIX</u>						
N 022020 001	7544370	Jun 07, 2026	DP			
	7550153	Sep 30, 2024		U-859		
	7553498	Sep 30, 2024		U-859		

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<u>PANTOPRAZOLE SODIUM - PROTONIX</u>						
N 022020	001 7838027	Sep 30, 2024	DP U-859			
<u>PARICALCITOL - ZEMPLAR</u>						
N 021606	001				ODE-125	Oct 18, 2023
<u>PARICALCITOL - ZEMPLAR</u>						
N 021606	002				ODE-125	Oct 18, 2023
<u>PARICALCITOL - ZEMPLAR</u>						
N 021606	003				ODE-125	Oct 18, 2023
<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		I-785	Jun 29, 2021
	7759308	Oct 25, 2026	DP		ODE-268	Jun 29, 2025
	8822637	Aug 06, 2023	U-1629		ODE-81	Dec 15, 2021
	9351923	May 23, 2028	DP			
<u>PASIREOTIDE PAMOATE - SIGNIFOR LAR KIT</u>						
N 203255	002 7473761	Dec 14, 2026	DS DP		I-785	Jun 29, 2021
	7759308	Oct 25, 2026	DP		ODE-268	Jun 29, 2025
	8822637	Aug 06, 2023	U-1629		ODE-81	Dec 15, 2021
	9351923	May 23, 2028	DP			
<u>PASIREOTIDE PAMOATE - SIGNIFOR LAR KIT</u>						
N 203255	003 7473761	Dec 14, 2026	DS DP		I-785	Jun 29, 2021
	7759308	Oct 25, 2026	DP		ODE-268	Jun 29, 2025
	8822637	Aug 06, 2023	U-1629		ODE-81	Dec 15, 2021
	9351923	May 23, 2028	DP			
<u>PASIREOTIDE PAMOATE - SIGNIFOR LAR KIT</u>						
N 203255	004 7473761	Dec 14, 2026	DS DP		I-785	Jun 29, 2021
	7759308	Oct 25, 2026	DP		ODE-268	Jun 29, 2025
	9351923	May 23, 2028	DP			
<u>PASIREOTIDE PAMOATE - SIGNIFOR LAR KIT</u>						
N 203255	005 7473761	Dec 14, 2026	DS DP		I-785	Jun 29, 2021
	7759308	Oct 25, 2026	DP		ODE-268	Jun 29, 2025
	9351923	May 23, 2028	DP			

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<u>PATIROMER SORBITEK CALCIUM - VELTASSA</u>						
N 205739 001	10485821	Mar 30, 2024	U-1766		NCE	Oct 21, 2020
	7556799	Feb 27, 2025	U-1766			
	8147873	Jun 20, 2028	DP			
	8216560	Mar 14, 2027	U-1766			
	8282913	May 29, 2027	DP			
	8287847	Mar 30, 2024	U-1766			
	8337824	May 29, 2030	DS U-1766			
	8475780	Mar 30, 2024	U-1766			
	8778324	Mar 30, 2024	U-1766			
	8889115	Mar 30, 2024	U-1766			
	9492476	Oct 08, 2033	U-1766			
	9925212	Oct 08, 2033	U-1766			
<u>PATIROMER SORBITEK CALCIUM - VELTASSA</u>						
N 205739 002	10485821	Mar 30, 2024	U-1766		NCE	Oct 21, 2020
	7556799	Feb 27, 2025	U-1766			
	8147873	Jun 20, 2028	DP			
	8216560	Mar 14, 2027	U-1766			
	8282913	May 29, 2027	DP			
	8287847	Mar 30, 2024	U-1766			
	8337824	May 29, 2030	DS U-1766			
	8475780	Mar 30, 2024	U-1766			
	8778324	Mar 30, 2024	U-1766			
	8889115	Mar 30, 2024	U-1766			
	9492476	Oct 08, 2033	U-1766			
	9925212	Oct 08, 2033	U-1766			
<u>PATIROMER SORBITEK CALCIUM - VELTASSA</u>						
N 205739 003	10485821	Mar 30, 2024	U-1766		NCE	Oct 21, 2020
	7556799	Feb 27, 2025	U-1766			
	8147873	Jun 20, 2028	DP			
	8216560	Mar 14, 2027	U-1766			
	8282913	May 29, 2027	DP			
	8287847	Mar 30, 2024	U-1766			
	8337824	May 29, 2030	DS U-1766			
	8475780	Mar 30, 2024	U-1766			
	8778324	Mar 30, 2024	U-1766			
	8889115	Mar 30, 2024	U-1766			
	9492476	Oct 08, 2033	U-1766			
	9925212	Oct 08, 2033	U-1766			
<u>PATISIRAN SODIUM - ONPATRO</u>						
N 210922 001	8058069	Apr 15, 2029	DP		NCE	Aug 10, 2023
	8158601	Nov 10, 2030	DP U-2378		ODE-197	Aug 10, 2025
	8168775	Oct 20, 2029	DS DP U-2378			
	8334373	May 27, 2025	DS DP			
	8362231	Mar 30, 2021	DS DP			
	8372968	Mar 30, 2021	DS DP			
	8492359	Apr 15, 2029	DP			
	8552171	Mar 30, 2021	DS DP			
	8642076	Oct 03, 2027	DP			
	8741866	Oct 20, 2029	U-2378			
	8778902	Mar 30, 2021	U-2378			
	8802644	Oct 21, 2030	DP U-2378			
	8822668	Apr 15, 2029	DP U-2378			
	8895718	Mar 30, 2021	DS DP			
	8895721	Mar 30, 2021	DS DP			
	9193753	Mar 30, 2021	U-2378			
	9234196	Oct 20, 2029	DP U-2378			
	9364435	Apr 15, 2029	DP U-2378			
	9567582	Mar 30, 2021	DS DP			
	9943538	Nov 04, 2023	DP			
	9943539	Nov 04, 2023	DP			
<u>PAZOPANIB HYDROCHLORIDE - VOTRIENT</u>						
N 022465 001	7105530	Oct 19, 2023	DS DP			
	7262203	Dec 19, 2021	DS DP			
	8114885	Dec 19, 2021	DS DP			

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<u>PAZOPANIB HYDROCHLORIDE - VOTRIENT</u>						
N 022465	002	7105530	Oct 19, 2023	DS DP		
		7262203	Dec 19, 2021	DS DP		
		8114885	Dec 19, 2021	DS DP		
<u>PEGINESATIDE ACETATE - OMONTYS PRESERVATIVE FREE</u>						
N 202799	001	7084245	May 12, 2024	DS DP	U-1238	
		7414105	May 12, 2024	DS DP	U-1238	
		7528104	May 12, 2024	DS DP		
		7550433	Jun 02, 2026		U-1238	
		7919118	May 12, 2024	DS DP		
		7919461	Jun 02, 2026		U-1238	
<u>PEGINESATIDE ACETATE - OMONTYS PRESERVATIVE FREE</u>						
N 202799	002	7084245	May 12, 2024	DS DP	U-1238	
		7414105	May 12, 2024	DS DP	U-1238	
		7528104	May 12, 2024	DS DP		
		7550433	Jun 02, 2026		U-1238	
		7919118	May 12, 2024	DS DP		
		7919461	Jun 02, 2026		U-1238	
<u>PEGINESATIDE ACETATE - OMONTYS PRESERVATIVE FREE</u>						
N 202799	003	7084245	May 12, 2024	DS DP	U-1238	
		7414105	May 12, 2024	DS DP	U-1238	
		7528104	May 12, 2024	DS DP		
		7550433	Jun 02, 2026		U-1238	
		7919118	May 12, 2024	DS DP		
		7919461	Jun 02, 2026		U-1238	
<u>PEGINESATIDE ACETATE - OMONTYS PRESERVATIVE FREE</u>						
N 202799	004	7084245	May 12, 2024	DS DP	U-1238	
		7414105	May 12, 2024	DS DP	U-1238	
		7528104	May 12, 2024	DS DP		
		7550433	Jun 02, 2026		U-1238	
		7919118	May 12, 2024	DS DP		
		7919461	Jun 02, 2026		U-1238	
<u>PEGINESATIDE ACETATE - OMONTYS PRESERVATIVE FREE</u>						
N 202799	005	7084245	May 12, 2024	DS DP	U-1238	
		7414105	May 12, 2024	DS DP	U-1238	
		7528104	May 12, 2024	DS DP		
		7550433	Jun 02, 2026		U-1238	
		7919118	May 12, 2024	DS DP		
		7919461	Jun 02, 2026		U-1238	
<u>PEGINESATIDE ACETATE - OMONTYS PRESERVATIVE FREE</u>						
N 202799	006	7084245	May 12, 2024	DS DP	U-1238	
		7414105	May 12, 2024	DS DP	U-1238	
		7528104	May 12, 2024	DS DP		
		7550433	Jun 02, 2026		U-1238	
		7919118	May 12, 2024	DS DP		
		7919461	Jun 02, 2026		U-1238	
<u>PEGINESATIDE ACETATE - OMONTYS</u>						
N 202799	007	7084245	May 12, 2024	DS DP	U-1238	
		7414105	May 12, 2024	DS DP	U-1238	
		7528104	May 12, 2024	DS DP		
		7550433	Jun 02, 2026		U-1238	
		7919118	May 12, 2024	DS DP		
		7919461	Jun 02, 2026		U-1238	
<u>PEGINESATIDE ACETATE - OMONTYS</u>						
N 202799	008	7084245	May 12, 2024	DS DP	U-1238	
		7414105	May 12, 2024	DS DP	U-1238	
		7528104	May 12, 2024	DS DP		
		7550433	Jun 02, 2026		U-1238	
		7919118	May 12, 2024	DS DP		
		7919461	Jun 02, 2026		U-1238	

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<u>PEMETREXED - PEMFEXY</u>						
N 209472	001	7772209	May 24, 2022	U-2728		
		7772209	May 24, 2022	U-2729		
<u>PEMETREXED DISODIUM - ALIMTA</u>						
N 021462	001	7772209	Nov 24, 2021	U-1296		
<u>PEMETREXED DISODIUM - ALIMTA</u>						
N 021462	002	7772209	Nov 24, 2021	U-1296		
<u>PENCICLOVIR - DENAVIR</u>						
N 020629	001	6579981	Jun 17, 2020	U-501		
<u>PERAMIVIR - RAPIVAB</u>						
N 206426	001	10391075	Feb 12, 2027	U-2622	NPP	Sep 20, 2020
		6562861	Dec 16, 2023	DS		
		8778997	May 07, 2027	U-1627		
		8778997	May 07, 2027	U-2622		
<u>PERAMPANEL - FYCOMPA</u>						
N 202834	001	6949571	Jun 08, 2021	DS DP U-106		
		6949571	Jun 08, 2021	DS DP U-2088		
		6949571	Jun 08, 2021	DS DP U-2089		
		6949571	Jun 08, 2021	DS DP U-2428		
		6949571	Jun 08, 2021	DS DP U-2429		
		8772497	Jul 01, 2026	DS		
<u>PERAMPANEL - FYCOMPA</u>						
N 202834	002	6949571	Jun 08, 2021	DS DP U-106		
		6949571	Jun 08, 2021	DS DP U-2088		
		6949571	Jun 08, 2021	DS DP U-2089		
		6949571	Jun 08, 2021	DS DP U-2428		
		6949571	Jun 08, 2021	DS DP U-2429		
		8772497	Jul 01, 2026	DS		
<u>PERAMPANEL - FYCOMPA</u>						
N 202834	003	6949571	Jun 08, 2021	DS DP U-106		
		6949571	Jun 08, 2021	DS DP U-2088		
		6949571	Jun 08, 2021	DS DP U-2089		
		6949571	Jun 08, 2021	DS DP U-2428		
		6949571	Jun 08, 2021	DS DP U-2429		
		8772497	Jul 01, 2026	DS		
<u>PERAMPANEL - FYCOMPA</u>						
N 202834	004	6949571	Jun 08, 2021	DS DP U-106		
		6949571	Jun 08, 2021	DS DP U-2088		
		6949571	Jun 08, 2021	DS DP U-2089		
		6949571	Jun 08, 2021	DS DP U-2428		
		6949571	Jun 08, 2021	DS DP U-2429		
		8772497	Jul 01, 2026	DS		
<u>PERAMPANEL - FYCOMPA</u>						
N 202834	005	6949571	Jun 08, 2021	DS DP U-106		
		6949571	Jun 08, 2021	DS DP U-2088		
		6949571	Jun 08, 2021	DS DP U-2089		
		6949571	Jun 08, 2021	DS DP U-2428		
		6949571	Jun 08, 2021	DS DP U-2429		
		8772497	Jul 01, 2026	DS		
<u>PERAMPANEL - FYCOMPA</u>						
N 202834	006	6949571	Jun 08, 2021	DS DP U-106		
		6949571	Jun 08, 2021	DS DP U-2088		
		6949571	Jun 08, 2021	DS DP U-2089		
		6949571	Jun 08, 2021	DS DP U-2428		
		6949571	Jun 08, 2021	DS DP U-2429		
		8772497	Jul 01, 2026	DS		
<u>PERAMPANEL - FYCOMPA</u>						
N 208277	001	6949571	Jun 08, 2021	DS DP U-106		
		6949571	Jun 08, 2021	DS DP U-2088		
		6949571	Jun 08, 2021	DS DP U-2089		

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<u>PERAMPANEL - FYCOMPA</u>						
N 208277	001	6949571	Jun 08, 2021	DS DP	U-2428	
		6949571	Jun 08, 2021	DS DP	U-2429	
		8772497	Jul 01, 2026	DS		
<u>PERFLUTREN - DEFINITY</u>						
N 021064	001	9789210	Mar 16, 2037		U-665	
<u>PEXIDARTINIB HYDROCHLORIDE - TURALIO</u>						
N 211810	001	10189833	May 05, 2036		U-2606	
		10435404	Jul 24, 2038	DP		NCE Aug 02, 2024
		7893075	Oct 13, 2028	DS		ODE-250 Aug 02, 2026
		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>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 - OSYMIA</u>						
N 022580	001	7056890	Jun 14, 2020	DP	U-1262	
		7553818	Jun 14, 2020		U-1262	
		7659256	Jun 14, 2020	DP	U-1262	
		7674776	Jun 14, 2020	DP	U-1262	
		8580298	May 15, 2029	DP		
		8580299	Jun 14, 2029		U-1262	
		8895057	Jun 09, 2028		U-1262	
		8895058	Jun 09, 2028	DP		
		9011905	Jun 09, 2028	DP		
		9011906	Jun 09, 2028		U-1262	
<u>PHENTERMINE HYDROCHLORIDE; TOPIRAMATE - OSYMIA</u>						
N 022580	002	7056890	Jun 14, 2020	DP	U-1262	
		7553818	Jun 14, 2020		U-1262	
		7659256	Jun 14, 2020	DP	U-1262	
		7674776	Jun 14, 2020	DP	U-1262	
		8580298	May 15, 2029	DP		
		8580299	Jun 14, 2029		U-1262	
		8895057	Jun 09, 2028		U-1262	
		8895058	Jun 09, 2028	DP		
		9011905	Jun 09, 2028	DP		
		9011906	Jun 09, 2028		U-1262	
<u>PHENTERMINE HYDROCHLORIDE; TOPIRAMATE - OSYMIA</u>						
N 022580	003	7056890	Jun 14, 2020	DP	U-1262	
		7553818	Jun 14, 2020		U-1262	
		7659256	Jun 14, 2020	DP	U-1262	
		7674776	Jun 14, 2020	DP	U-1262	
		8580298	May 15, 2029	DP		
		8580299	Jun 14, 2029		U-1262	
		8895057	Jun 09, 2028		U-1262	
		8895058	Jun 09, 2028	DP		
		9011905	Jun 09, 2028	DP		
		9011906	Jun 09, 2028		U-1262	
<u>PHENTERMINE HYDROCHLORIDE; TOPIRAMATE - OSYMIA</u>						
N 022580	004	7056890	Jun 14, 2020	DP	U-1262	
		7553818	Jun 14, 2020		U-1262	
		7659256	Jun 14, 2020	DP	U-1262	
		7674776	Jun 14, 2020	DP	U-1262	
		8580298	May 15, 2029	DP		
		8580299	Jun 14, 2029		U-1262	

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<u>PHENTERMINE HYDROCHLORIDE; TOPIRAMATE - QSYMIA</u>						
N 022580 004	8895057	Jun 09, 2028	U-1262			
	8895058	Jun 09, 2028	DP			
	9011905	Jun 09, 2028	DP			
	9011906	Jun 09, 2028	U-1262			
<u>PHENTOLAMINE MESYLATE - ORAVERSE</u>						
N 022159 001	6764678	May 11, 2021	U-967			
	6872390	May 11, 2021	DP			
	7229630	Jun 20, 2023	DP			
	7569230	Oct 17, 2023	U-967			
	7575757	Apr 21, 2025	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>PIMAVANSERIN TARTRATE - NUPLAZID</u>						
N 207318 001	10028944	Jan 15, 2024	U-1974		NCE	Apr 29, 2021
	6756393	Mar 06, 2021	DS DP			
	6815458	Mar 06, 2021	DS DP		U-1843	
	7115634	Oct 06, 2021	DS DP			
	7601740	Jun 17, 2027	DS DP			
	7659285	Aug 24, 2026			U-1844	
	7732615	Jun 03, 2028	DS DP			
	7858789	Dec 13, 2020	DS DP			
	7923564	Sep 26, 2025	DS DP			
	8110574	Dec 13, 2020	DS DP			
	8618130	Jan 15, 2024			U-1845	
	8921393	Jan 15, 2024			U-1846	
	9296694	Mar 06, 2021	DS DP			
	9566271	Jan 15, 2024			U-1974	
	9765053	Jul 27, 2022			U-1974	
<u>PIMAVANSERIN TARTRATE - NUPLAZID</u>						
N 207318 002	10028944	Jan 15, 2024	U-1974		NCE	Apr 29, 2021
	10517860	Mar 23, 2037	U-1974			
	6756393	Mar 06, 2021	DS DP			
	6815458	Mar 06, 2021	DS DP		U-1843	
	7115634	Oct 06, 2021	DS DP			
	7601740	Jun 17, 2027	DS DP			
	7659285	Aug 24, 2026			U-1844	
	7732615	Jun 03, 2028	DS DP			
	7858789	Dec 13, 2020	DS DP			
	7923564	Sep 26, 2025	DS DP			
	8110574	Dec 13, 2020	DS DP			
	8618130	Jan 15, 2024			U-1845	
	8921393	Jan 15, 2024			U-1846	
	9296694	Mar 06, 2021	DS DP			
	9566271	Jan 15, 2024			U-1974	
	9765053	Jul 27, 2022			U-1974	
<u>PIMAVANSERIN TARTRATE - NUPLAZID</u>						
N 210793 001	10028944	Jan 15, 2024	U-1974		NCE	Apr 29, 2021
	10449185	Aug 27, 2038	DP			
	6756393	Mar 06, 2021	DS DP			
	6815458	Mar 06, 2021	DS DP		U-1843	
	7115634	Oct 06, 2021	DS DP			
	7601740	Jun 17, 2027	DS DP			
	7659285	Aug 24, 2026			U-1844	
	7732615	Jun 03, 2028	DS DP			
	7858789	Dec 13, 2020	DS DP			
	7923564	Sep 26, 2025	DS DP			
	8110574	Dec 13, 2020	DS DP			
	8618130	Jan 15, 2024			U-1845	
	8921393	Jan 15, 2024			U-1846	
	9296694	Mar 06, 2021	DS DP			
	9566271	Jan 15, 2024			U-1974	
	9765053	Jul 27, 2022			U-1974	

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<u>PIMAVANSERIN TARTRATE - NUPLAZID</u>						
N 210793 001	10028944	Jan 15, 2024	U-1974		NCE	Apr 29, 2021
	10449185	Aug 27, 2038	DP			
	6756393	Mar 06, 2021	DS DP			
	6815458	Mar 06, 2021	DS DP	U-1843		
	7115634	Oct 06, 2021	DS DP			
	7601740	Jun 17, 2027	DS DP			
	7659285	Aug 24, 2026	U-1844			
	7732615	Jun 03, 2028	DS DP			
	7858789	Dec 13, 2020	DS DP			
	7923564	Sep 26, 2025	DS DP			
	8110574	Dec 13, 2020	DS DP			
	8618130	Jan 15, 2024	U-1845			
	8921393	Jan 15, 2024	U-1846			
	9296694	Mar 06, 2021	DS DP			
	9566271	Jan 15, 2024	U-1974			
	9765053	Jul 27, 2022	U-1974			
<u>PIPERACILLIN SODIUM; TAZOBACTAM SODIUM - ZOSYN</u>						
N 050684 001	6900184	Apr 14, 2023	DP U-282			
	7915229	Apr 14, 2023	DP			
	8133883	Apr 14, 2023	DP U-282			
<u>PIPERACILLIN SODIUM; TAZOBACTAM SODIUM - ZOSYN</u>						
N 050684 002	6900184	Apr 14, 2023	DP U-282			
	7915229	Apr 14, 2023	DP			
	8133883	Apr 14, 2023	DP U-282			
<u>PIPERACILLIN SODIUM; TAZOBACTAM SODIUM - ZOSYN</u>						
N 050684 003	6900184	Apr 14, 2023	DP U-282			
	7915229	Apr 14, 2023	DP			
	8133883	Apr 14, 2023	DP U-282			
<u>PIPERACILLIN SODIUM; TAZOBACTAM SODIUM - ZOSYN</u>						
N 050684 004	6900184	Apr 14, 2023	DP U-282			
	7915229	Apr 14, 2023	DP			
	8133883	Apr 14, 2023	DP U-282			
<u>PIPERACILLIN SODIUM; TAZOBACTAM SODIUM - ZOSYN IN PLASTIC CONTAINER</u>						
N 050750 001	6900184	Apr 14, 2023	DP U-282			
	7915229	Apr 14, 2023	DP			
	8133883	Apr 14, 2023	DP U-282			
<u>PIPERACILLIN SODIUM; TAZOBACTAM SODIUM - ZOSYN IN PLASTIC CONTAINER</u>						
N 050750 002	6900184	Apr 14, 2023	DP U-282			
	7915229	Apr 14, 2023	DP			
	8133883	Apr 14, 2023	DP U-282			
<u>PIPERACILLIN SODIUM; TAZOBACTAM SODIUM - ZOSYN IN PLASTIC CONTAINER</u>						
N 050750 003	6900184	Apr 14, 2023	DP U-282			
	7915229	Apr 14, 2023	DP			
	8133883	Apr 14, 2023	DP U-282			
<u>PIRFENIDONE - ESBRIET</u>						
N 022535 001	7566729	Apr 22, 2029	U-1600		ODE-77	Oct 15, 2021
	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			
	8383150	Sep 22, 2026	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			

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<u>PIRFENIDONE - ESBRIET</u>						
N 022535	001	8754109				
		8778947		U-1612		
				U-1613		
<u>PIRFENIDONE - ESBRIET</u>						
N 208780	001	10188637		DP	ODE-77	Oct 15, 2021
		7566729		U-2077		
		7566729		U-2078		
		7635707		U-2072		
		7635707		U-2073		
		7635707		U-2074		
		7635707		U-2075		
		7635707		U-2076		
		7635707		U-2083		
		7767700		U-2080		
		7816383		U-2042		
		7816383		U-2050		
		7910610		U-2048		
		7910610		U-2049		
		8013002		U-2047		
		8013002		U-2082		
		8084475		U-2052		
		8084475		U-2054		
		8318780		U-2046		
		8318780		U-2081		
		8383150		DP U-2361		
		8420674		U-2079		
		8592462		U-2055		
		8592462		U-2056		
		8592462		U-2057		
		8592462		U-2058		
		8592462		U-2059		
		8592462		U-2060		
		8592462		U-2061		
		8592462		U-2062		
		8592462		U-2063		
		8609701		U-2064		
		8609701		U-2065		
		8609701		U-2066		
		8609701		U-2067		
		8609701		U-2068		
		8609701		U-2069		
		8609701		U-2070		
		8648098		U-2051		
		8648098		U-2052		
		8754109		U-2053		
		8778947		U-2044		
		8778947		U-2045		
		9561217		DP		
<u>PIRFENIDONE - ESBRIET</u>						
N 208780	002	10188637		DP	ODE-77	Oct 15, 2021
		7566729		U-2269		
		7566729		U-2270		
		7635707		U-2072		
		7635707		U-2073		
		7635707		U-2074		
		7635707		U-2075		
		7635707		U-2076		
		7635707		U-2083		
		7767700		U-2080		
		7816383		U-2042		
		7816383		U-2050		
		7910610		U-2048		
		7910610		U-2049		
		8013002		U-2047		
		8013002		U-2082		
		8084475		U-2054		
		8084475		U-2268		
		8318780		U-2046		

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<u>PIRFENIDONE - ESBRIET</u>						
N 208780 002	8318780	Jan 08, 2030				
	8383150	Sep 22, 2026	DP			
	8420674	Dec 18, 2027				
	8592462	Apr 22, 2029				
	8592462	Apr 22, 2029				
	8592462	Apr 22, 2029				
	8592462	Apr 22, 2029				
	8592462	Apr 22, 2029				
	8592462	Apr 22, 2029				
	8592462	Apr 22, 2029				
	8592462	Apr 22, 2029				
	8592462	Apr 22, 2029				
	8609701	Apr 22, 2029				
	8609701	Apr 22, 2029				
	8609701	Apr 22, 2029				
	8609701	Apr 22, 2029				
	8609701	Apr 22, 2029				
	8609701	Apr 22, 2029				
	8609701	Apr 22, 2029				
	8609701	Apr 22, 2029				
	8609701	Apr 22, 2029				
	8609701	Apr 22, 2029				
	8609701	Apr 22, 2029				
	8609701	Apr 22, 2029				
	8648098	Jan 08, 2030				
	8648098	Jan 08, 2030				
	8754109	Jan 08, 2030				
	8778947	Aug 30, 2033				
	8778947	Aug 30, 2033				
	9561217	Jan 25, 2022	DP			
<u>PIRFENIDONE - ESBRIET</u>						
N 208780 003	10188637	Mar 28, 2037	DP		ODE-77	Oct 15, 2021
	7566729	Apr 22, 2029				
	7566729	Apr 22, 2029				
	7635707	Apr 22, 2029				
	7635707	Apr 22, 2029				
	7635707	Apr 22, 2029				
	7635707	Apr 22, 2029				
	7635707	Apr 22, 2029				
	7635707	Apr 22, 2029				
	7635707	Apr 22, 2029				
	7767700	Dec 18, 2027				
	7816383	Jan 08, 2030				
	7816383	Jan 08, 2030				
	7910610	Jan 08, 2030				
	7910610	Jan 08, 2030				
	8013002	Jan 08, 2030				
	8013002	Jan 08, 2030				
	8084475	Jan 08, 2030				
	8084475	Jan 08, 2030				
	8318780	Jan 08, 2030				
	8318780	Jan 08, 2030				
	8383150	Sep 22, 2026	DP			
	8420674	Dec 18, 2027				
	8592462	Apr 22, 2029				
	8592462	Apr 22, 2029				
	8592462	Apr 22, 2029				
	8592462	Apr 22, 2029				
	8592462	Apr 22, 2029				
	8592462	Apr 22, 2029				
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	8592462	Apr 22, 2029				
	8592462	Apr 22, 2029				
	8592462	Apr 22, 2029				
	8592462	Apr 22, 2029				
	8592462	Apr 22, 2029				
	8609701	Apr 22, 2029				
	8609701	Apr 22, 2029				
	8609701	Apr 22, 2029				
	8609701	Apr 22, 2029				
	8609701	Apr 22, 2029				
	8609701	Apr 22, 2029				
	8609701	Apr 22, 2029				
	8609701	Apr 22, 2029				
	8609701	Apr 22, 2029				
	8609701	Apr 22, 2029				
	8609701	Apr 22, 2029				
	8648098	Jan 08, 2030				
	8648098	Jan 08, 2030				
	8754109	Jan 08, 2030				
	8778947	Aug 30, 2033				

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<u>PIRFENIDONE - ESBRIET</u>						
N 208780 003	8778947	Aug 30, 2033				
	9561217	Jan 25, 2022	U-2045 DP			
<u>PITAVASTATIN CALCIUM - LIVALO</u>						
N 022363 001	5856336	Dec 25, 2020	DS	U-998	NPP	May 16, 2022
	5856336*PED	Jun 25, 2021			PED	Nov 16, 2022
	7022713	Feb 19, 2024		U-998		
	7022713*PED	Aug 19, 2024				
	8557993	Feb 02, 2024	DP			
	8557993*PED	Aug 02, 2024				
<u>PITAVASTATIN CALCIUM - LIVALO</u>						
N 022363 002	5856336	Dec 25, 2020	DS	U-998	NPP	May 16, 2022
	5856336*PED	Jun 25, 2021			PED	Nov 16, 2022
	7022713	Feb 19, 2024		U-998		
	7022713*PED	Aug 19, 2024				
	8557993	Feb 02, 2024	DP			
	8557993*PED	Aug 02, 2024				
<u>PITAVASTATIN CALCIUM - LIVALO</u>						
N 022363 003	5856336	Dec 25, 2020	DS	U-998	NPP	May 16, 2022
	5856336*PED	Jun 25, 2021			PED	Nov 16, 2022
	7022713	Feb 19, 2024		U-998		
	7022713*PED	Aug 19, 2024				
	8557993	Feb 02, 2024	DP			
	8557993*PED	Aug 02, 2024				
<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	7169928	Feb 02, 2020	DS DP		NCE	Aug 14, 2024
	7910605	Sep 23, 2022		U-1101	ODE-255	Aug 14, 2026
	8207197	Feb 25, 2029	DS DP			
	8354430	Feb 26, 2026		U-1101		
	8486947	Sep 26, 2029		U-1101		
<u>PITOLISANT HYDROCHLORIDE - WAKIX</u>						
N 211150 002	7169928	Feb 02, 2020	DS DP		NCE	Aug 14, 2024
	7910605	Sep 23, 2022		U-1101	ODE-255	Aug 14, 2026
	8207197	Feb 25, 2029	DS DP			
	8354430	Feb 26, 2026		U-1101		
	8486947	Sep 26, 2029		U-1101		
<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		I-764	Jan 24, 2021
	7041786	Mar 25, 2023	DS		NCE	Jan 19, 2022
	7799897	Jun 09, 2022	DS			
	8637451	Mar 28, 2022		U-1964		
	9610321	Sep 15, 2031		U-1999		
	9610321	Sep 15, 2031		U-2230		
	9616097	Jul 02, 2032	DP			
	9919024	Sep 15, 2031		U-1999		
	9919024	Sep 15, 2031		U-2230		
	9925231	Sep 15, 2031	DP			

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<u>PLERIXAFOR - MOZOBIL</u>						
N 022311	001	6987102	Jul 22, 2023	U-936		
		7897590	Jul 22, 2023	U-936		
<u>POLIDOCANOL - VARITHENA</u>						
N 205098	001	6572873	May 26, 2020	U-1461		
		6846412	Jul 19, 2022	DP		
		6942165	May 26, 2020	DP		
		7025290	May 26, 2020	DP	U-1461	
		7357336	May 26, 2020	U-1461		
		7604185	May 26, 2020	DS DP	U-1462	
		7731986	Nov 17, 2024	DS DP	U-1463	
		7814943	Nov 19, 2027	DP	U-1461	
		7842282	May 26, 2020	U-1461		
		7842283	May 26, 2020	DP		
		8122917	Sep 09, 2024	DP		
		8323677	May 26, 2020	DS		
		8734833	May 26, 2020	DS DP		
		9480652	May 12, 2032	DP		
<u>POMALIDOMIDE - POMALYST</u>						
N 204026	001	10555939	May 19, 2030	DP	ODE-43	Feb 08, 2020
		6315720	Oct 23, 2020	U-1361		
		6561977	Oct 23, 2020	U-1361		
		6755784	Oct 23, 2020	U-1361		
		8198262	Jun 17, 2025	U-1360		
		8198262	Jun 17, 2025	U-2254		
		8315886	Oct 23, 2020	U-1361		
		8626531	Oct 23, 2020	U-1361		
		8673939	May 15, 2023	U-1360		
		8673939	May 15, 2023	U-2254		
		8735428	May 15, 2023	U-1360		
		8735428	May 15, 2023	U-2254		
		8828427	Jun 21, 2031	DS DP		
		9993467	May 19, 2030	DP		
<u>POMALIDOMIDE - POMALYST</u>						
N 204026	002	10555939	May 19, 2030	DP	ODE-43	Feb 08, 2020
		6315720	Oct 23, 2020	U-1361		
		6561977	Oct 23, 2020	U-1361		
		6755784	Oct 23, 2020	U-1361		
		8198262	Jun 17, 2025	U-1360		
		8198262	Jun 17, 2025	U-2254		
		8315886	Oct 23, 2020	U-1361		
		8626531	Oct 23, 2020	U-1361		
		8673939	May 15, 2023	U-1360		
		8673939	May 15, 2023	U-2254		
		8735428	May 15, 2023	U-1360		
		8735428	May 15, 2023	U-2254		
		8828427	Jun 21, 2031	DS DP		
		9993467	May 19, 2030	DP		
<u>POMALIDOMIDE - POMALYST</u>						
N 204026	003	10555939	May 19, 2030	DP	ODE-43	Feb 08, 2020
		6315720	Oct 23, 2020	U-1361		
		6561977	Oct 23, 2020	U-1361		
		6755784	Oct 23, 2020	U-1361		
		8198262	Jun 17, 2025	U-1360		
		8198262	Jun 17, 2025	U-2254		
		8315886	Oct 23, 2020	U-1361		
		8626531	Oct 23, 2020	U-1361		
		8673939	May 15, 2023	U-1360		
		8673939	May 15, 2023	U-2254		
		8735428	May 15, 2023	U-1360		
		8735428	May 15, 2023	U-2254		
		8828427	Jun 21, 2031	DS DP		
		9993467	May 19, 2030	DP		

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<u>POMALIDOMIDE - POMALYST</u>						
N 204026 004	10555939	May 19, 2030	DP		ODE-43	Feb 08, 2020
	6315720	Oct 23, 2020	U-1361			
	6561977	Oct 23, 2020	U-1361			
	6755784	Oct 23, 2020	U-1361			
	8198262	Jun 17, 2025	U-1360			
	8198262	Jun 17, 2025	U-2254			
	8315886	Oct 23, 2020	U-1361			
	8626531	Oct 23, 2020	U-1361			
	8673939	May 15, 2023	U-1360			
	8673939	May 15, 2023	U-2254			
	8735428	May 15, 2023	U-1360			
	8735428	May 15, 2023	U-2254			
	8828427	Jun 21, 2031	DS DP			
	9993467	May 19, 2030	DP			
<u>PONATINIB HYDROCHLORIDE - ICLUSIG</u>						
N 203469 001	8114874	Dec 22, 2026	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	8114874	Dec 22, 2026	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	8114874	Dec 22, 2026	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>POSACONAZOLE - NOXAFIL</u>						
N 022003 001	8263600	Apr 01, 2022	DP			
<u>POSACONAZOLE - NOXAFIL</u>						
N 205596 001	10117951	Mar 13, 2029	DP			
	8410077	Mar 13, 2029	DP			
	9023790	Jul 04, 2031	DP	U-1698		
	9358297	Jun 24, 2031	DP	U-1454		
	9493582	Feb 27, 2033	DP			
	9750822	Mar 13, 2029	DP			
<u>PRALATREXATE - FOLOTYN</u>						
N 022468 001	6028071	Jul 16, 2022	DS DP	U-1004		
	7622470	May 31, 2025		U-1015		
	8299078	May 31, 2025		U-1004		
<u>PRALATREXATE - FOLOTYN</u>						
N 022468 002	6028071	Jul 16, 2022	DS DP	U-1004		
	7622470	May 31, 2025		U-1015		
	8299078	May 31, 2025		U-1004		
<u>PRAMIPEXOLE DIHYDROCHLORIDE - MIRAPEX ER</u>						
N 022421 001	7695734	Apr 26, 2028	DP			
	8679533	Sep 08, 2029	DP	U-219		

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<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			
<u>PRASTERONE - INTRAROSA</u>						
N 208470 001	8268806	Mar 19, 2031	DP		NCE	Nov 16, 2021
	8629129	Aug 07, 2028	DP			
	8957054	Aug 07, 2028	U-1922			
<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	6488960	Mar 14, 2020	DP U-1267			
	6677326	Mar 14, 2020	DP U-1268			
	8309124	Apr 23, 2024	U-1292			
	8394407	Apr 23, 2024	DP U-1362			
	9040085	Apr 23, 2024	U-1362			
	9186332	Apr 23, 2024	U-1362			
	9504699	Aug 03, 2027	U-1362			
<u>PREDNISONE - RAYOS</u>						
N 202020 002	6488960	Mar 14, 2020	DP U-1267			
	6677326	Mar 14, 2020	DP U-1268			
	8309124	Apr 23, 2024				
	8394407	Apr 23, 2024	DP U-1362			
	9040085	Apr 23, 2024	U-1362			
	9186332	Apr 23, 2024	U-1362			
	9504699	Aug 03, 2027	U-1362			
<u>PREDNISONE - RAYOS</u>						
N 202020 003	8168218	Jan 07, 2028	DP U-1269			
	8309124	Apr 23, 2024	U-1292			
	8394407	Apr 23, 2024	DP U-1362			
	9040085	Apr 23, 2024	U-1362			
	9186332	Apr 23, 2024	U-1362			
	9504699	Aug 03, 2027	U-1362			
<u>PREGABALIN - LYRICA</u>						
N 021446 001					NPP	May 03, 2021
					NPP	May 23, 2022
					PED	Nov 03, 2021
					PED	Nov 23, 2022

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<u>PREGABALIN - LYRICA</u>						
N 021446	002				NPP	May 03, 2021
					NPP	May 23, 2022
					PED	Nov 03, 2021
					PED	Nov 23, 2022
<u>PREGABALIN - LYRICA</u>						
N 021446	003				NPP	May 03, 2021
					NPP	May 23, 2022
					PED	Nov 03, 2021
					PED	Nov 23, 2022
<u>PREGABALIN - LYRICA</u>						
N 021446	004				NPP	May 03, 2021
					NPP	May 23, 2022
					PED	Nov 03, 2021
					PED	Nov 23, 2022
<u>PREGABALIN - LYRICA</u>						
N 021446	005				NPP	May 03, 2021
					NPP	May 23, 2022
					PED	Nov 03, 2021
					PED	Nov 23, 2022
<u>PREGABALIN - LYRICA</u>						
N 021446	006				NPP	May 03, 2021
					NPP	May 23, 2022
					PED	Nov 03, 2021
					PED	Nov 23, 2022
<u>PREGABALIN - LYRICA</u>						
N 021446	007				NPP	May 03, 2021
					NPP	May 23, 2022
					PED	Nov 03, 2021
					PED	Nov 23, 2022
<u>PREGABALIN - LYRICA</u>						
N 021446	008				NPP	May 03, 2021
					NPP	May 23, 2022
					PED	Nov 03, 2021
					PED	Nov 23, 2022
<u>PREGABALIN - LYRICA</u>						
N 022488	001				NPP	May 03, 2021
					NPP	May 23, 2022
					PED	Nov 03, 2021
					PED	Nov 23, 2022
<u>PREGABALIN - LYRICA CR</u>						
N 209501	001	10022447	Nov 02, 2026	U-2136	NP	Oct 11, 2020
		10022447	Nov 02, 2026	U-2137	PED	Apr 11, 2021
		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	NP	Oct 11, 2020
		10022447	Nov 02, 2026	U-2137	PED	Apr 11, 2021
		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	NP	Oct 11, 2020
		10022447	Nov 02, 2026	U-2137	PED	Apr 11, 2021

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<u>PREGABALIN - LYRICA CR</u>						
N 209501 003	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>PRETOMANID - PRETOMANID</u>						
N 212862 001					NCE	Aug 14, 2024
					ODE-253	Aug 14, 2026
					GAIN	Aug 14, 2029
<u>PROPOFOL - DIPRIVAN</u>						
N 019627 002	8476010	Dec 01, 2024	DS DP			
	8476010*PED	Jun 01, 2025				
<u>PROPRANOLOL HYDROCHLORIDE - INNOPRAN XL</u>						
N 021438 001	6500454	Oct 04, 2021	DP			
<u>PROPRANOLOL HYDROCHLORIDE - INNOPRAN XL</u>						
N 021438 002	6500454	Oct 04, 2021	DP			
<u>PROPRANOLOL HYDROCHLORIDE - HEMANGEOL</u>						
N 205410 001	8338489	Oct 16, 2028		U-1496	ODE-62	Mar 14, 2021
	8987262	Oct 16, 2028		U-1988		
<u>PRUCALOPRIDE SUCCINATE - MOTEGRITY</u>						
N 210166 001					NCE	Dec 14, 2023
<u>PRUCALOPRIDE SUCCINATE - MOTEGRITY</u>						
N 210166 002					NCE	Dec 14, 2023
<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>RADIUM RA-223 DICHLORIDE - XOFIGO</u>						
N 203971 001	6635234	Nov 17, 2022		U-2271		
<u>RALTEGRAVIR POTASSIUM - ISENTRESS</u>						
N 022145 001	7169780	Oct 03, 2023	DS DP		D-167	May 26, 2020
	7169780*PED	Apr 03, 2024			NPP	Nov 22, 2020
	7217713	Oct 21, 2022		U-257	PED	Nov 26, 2020
	7217713*PED	Apr 21, 2023			PED	May 22, 2021
	7435734	Oct 21, 2022		U-257		
	7435734	Oct 21, 2022		U-900		
	7435734*PED	Apr 21, 2023				
	7754731	Mar 11, 2029	DS DP	U-257		
	7754731*PED	Sep 11, 2029				
<u>RALTEGRAVIR POTASSIUM - ISENTRESS HD</u>						
N 022145 002	7169780	Oct 03, 2023	DS DP		NPP	Nov 22, 2020
	7169780*PED	Apr 03, 2024			NS	May 26, 2020
	7217713	Oct 21, 2022		U-257	PED	Nov 26, 2020
	7217713*PED	Apr 21, 2023			PED	May 22, 2021
	7435734	Oct 21, 2022		U-257		
	7435734	Oct 21, 2022		U-900		
	7435734*PED	Apr 21, 2023				
	7754731	Mar 11, 2029	DS DP	U-257		
	7754731*PED	Sep 11, 2029				
	9649311	Oct 21, 2030		DP		
	9649311*PED	Apr 21, 2031				
<u>RALTEGRAVIR POTASSIUM - ISENTRESS</u>						
N 203045 001	7169780	Oct 03, 2023	DS DP		NPP	Nov 22, 2020
	7169780*PED	Apr 03, 2024			PED	May 22, 2021
	7217713	Oct 21, 2022		U-257		
	7217713*PED	Apr 21, 2023				
	7435734	Oct 21, 2022		U-257		

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<u>RALTEGRAVIR POTASSIUM - ISENTRESS</u>						
N 203045	001	7435734*PED	Apr 21, 2023			
		7754731	Mar 11, 2029	DS DP	U-257	
		7754731*PED	Sep 11, 2029			
<u>RALTEGRAVIR POTASSIUM - ISENTRESS</u>						
N 203045	002	7169780	Oct 03, 2023	DS DP		NPP Nov 22, 2020
		7169780*PED	Apr 03, 2024			PED May 22, 2021
		7217713	Oct 21, 2022		U-257	
		7217713*PED	Apr 21, 2023			
		7435734	Oct 21, 2022		U-257	
		7435734*PED	Apr 21, 2023			
		7754731	Mar 11, 2029	DS DP	U-257	
		7754731*PED	Sep 11, 2029			
<u>RALTEGRAVIR POTASSIUM - ISENTRESS</u>						
N 205786	001	7169780	Oct 03, 2023	DS DP		NPP Nov 22, 2020
		7169780*PED	Apr 03, 2024			PED May 22, 2021
		7217713	Oct 21, 2022		U-257	
		7217713*PED	Apr 21, 2023			
		7435734	Oct 21, 2022		U-257	
		7435734*PED	Apr 21, 2023			
		7754731	Mar 11, 2029	DS DP	U-257	
		7754731*PED	Sep 11, 2029			
<u>RAMELTEON - ROZEREM</u>						
N 021782	001	10098866	Nov 16, 2021	DP	U-2433	
<u>RAMIPRIL - ALTACE</u>						
N 019901	001	7368469	Aug 30, 2020		U-871	
<u>RAMIPRIL - ALTACE</u>						
N 019901	002	7368469	Aug 30, 2020		U-871	
<u>RAMIPRIL - ALTACE</u>						
N 019901	003	7368469	Aug 30, 2020		U-871	
<u>RAMIPRIL - ALTACE</u>						
N 019901	004	7368469	Aug 30, 2020		U-871	
<u>RAMIPRIL - ALTACE</u>						
N 022021	001	7368469	Aug 30, 2020		U-871	
<u>RAMIPRIL - ALTACE</u>						
N 022021	002	7368469	Aug 30, 2020		U-871	
<u>RAMIPRIL - ALTACE</u>						
N 022021	003	7368469	Aug 30, 2020		U-871	
<u>RAMIPRIL - ALTACE</u>						
N 022021	004	7368469	Aug 30, 2020		U-871	
<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	6403567	Apr 10, 2022	DS DP	U-869	M-194 Jan 17, 2020
		8106183	Feb 02, 2027	DS		
		RE47301	Feb 02, 2027		DP	
<u>REGORAFENIB - STIVARGA</u>						
N 203085	001	7351834	Jun 28, 2022	DS		I-744 Apr 27, 2020
		8637553	Feb 16, 2031	DS DP		ODE-139 Apr 27, 2024
		8680124	Jun 02, 2030		U-1506	ODE-44 Feb 25, 2020
		9458107	Apr 08, 2031		DP	
		9957232	Jul 09, 2032	DS		

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<u>RETAPAMULIN - ALTABAX</u>						
N 022055	001 7875630	Feb 14, 2027	DS			
	8207191	Aug 30, 2024		U-805		
	RE43390	Apr 12, 2021	DS DP	U-805		
<u>REVEFENACIN - YUPELRI</u>						
N 210598	001 10106503	Mar 10, 2025		U-2440	NCE	Nov 09, 2023
	10343995	Mar 10, 2025		U-2440		
	10550081	Jul 14, 2030	DS			
	7288657	Dec 23, 2025	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>RIBAVIRIN - REBETOL</u>						
N 021546	001 6790837	Apr 05, 2023		DP		
<u>RIBOCICLIB SUCCINATE - KISQALI</u>						
N 209092	001 8324225	Jun 17, 2028	DS DP		I-783	Jul 18, 2021
	8415355	Feb 19, 2031	DS DP		I-784	Jul 18, 2021
	8685980	May 25, 2030	DS DP		NCE	Mar 13, 2022
	8962630	Dec 09, 2029		U-1981		
	8962630	Dec 09, 2029		U-2355		
	8962630	Dec 09, 2029		U-2356		
	9193732	Nov 09, 2031	DS DP			
	9416136	Aug 20, 2029		U-1981		
	9416136	Aug 20, 2029		U-2355		
	9416136	Aug 20, 2029		U-2356		
	9868739	Nov 09, 2031		U-1981		
	9868739	Nov 09, 2031		U-2355		
	9868739	Nov 09, 2031		U-2356		
<u>RIBOFLAVIN 5'-PHOSPHATE SODIUM - PHOTREXA</u>						
N 203324	001				ODE-116	Apr 15, 2023
					ODE-121	Jul 15, 2023
<u>RIBOFLAVIN 5'-PHOSPHATE SODIUM - PHOTREXA VISCOUS IN DEXTRAN 20%</u>						
N 203324	002				ODE-116	Apr 15, 2023
					ODE-121	Jul 15, 2023
<u>RIFAMYCIN SODIUM - AEMCOLO</u>						
N 210910	001 8263120	May 03, 2025		DP		
	8486446	May 03, 2025		DP		
	8529945	May 03, 2025		DP		
	8741948	May 03, 2025		DP	U-2448	
<u>RIFAXIMIN - XIFAXAN</u>						
N 021361	001 7045620	Jun 19, 2024	DS DP			
	7612199	Jun 19, 2024	DS DP			
	7902206	Jun 19, 2024	DS DP			
	7906542	Jun 01, 2025	DS DP			
	7928115	Jul 24, 2029		U-1121		
	8158644	Jun 19, 2024		DP		
	8158781	Jun 19, 2024	DS			
	8193196	Sep 02, 2027	DS DP			
	8518949	Feb 27, 2026		DP		
	8741904	Feb 27, 2026	DS	U-1526		
	8835452	Jun 19, 2024	DS DP			
	8853231	Jun 19, 2024		DP		
	9271968	Feb 27, 2026		DP		
<u>RIFAXIMIN - XIFAXAN</u>						
N 022554	001 10314828	Oct 02, 2029		U-1481		
	10335397	Oct 02, 2029		U-2579		

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APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>RIFAXIMIN - XIFAXAN</u>						
N 022554	001	10456384	Feb 26, 2029	U-2643		
		10456384	Feb 26, 2029	U-2644		
		7045620	Jun 19, 2024	DS		
		7612199	Jun 19, 2024	DS DP		
		7902206	Jun 19, 2024	DS DP		
		7906542	Jun 01, 2025	DS DP		
		7915275	Feb 23, 2025		U-1707	
		7915275	Feb 23, 2025		U-1708	
		8158644	Jun 19, 2024	DP		
		8158781	Jun 19, 2024	DS		
		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	
		8835452	Jun 19, 2024	DS DP		
		8853231	Jun 19, 2024	DP		
		8946252	Jul 24, 2029		U-1481	
		8969398	Oct 02, 2029		U-1481	
		9271968	Feb 27, 2026	DP		
		9421195	Mar 10, 2030		U-1481	
		9629828	Jul 24, 2029		U-1994	
<u>RILPIVIRINE HYDROCHLORIDE - EDURANT</u>						
N 202022	001	6838464	Feb 26, 2021	DS DP	M-223	Feb 01, 2021
		7125879	Apr 21, 2025	DS DP	U-1153	
		7125879	Apr 21, 2025	DS DP	U-1307	
		7125879	Apr 21, 2025	DS DP	U-1740	
		7638522	Apr 14, 2023	DP		
		8080551	Apr 11, 2023	DS DP		
		8101629	Aug 09, 2022	DP		
<u>RILUZOLE - TIGLUTIK KIT</u>						
N 209080	001	8765150	Mar 12, 2029	DP	U-2401	
<u>RIMEGEPANT SULFATE - NURTEC ODT</u>						
N 212728	001				NCE	Feb 27, 2025
<u>RIOCIGUAT - ADEMPAS</u>						
N 204819	001	7173037	Dec 04, 2026	DS DP	ODE-53	Oct 08, 2020
<u>RIOCIGUAT - ADEMPAS</u>						
N 204819	002	7173037	Dec 04, 2026	DS DP	ODE-53	Oct 08, 2020
<u>RIOCIGUAT - ADEMPAS</u>						
N 204819	003	7173037	Dec 04, 2026	DS DP	ODE-53	Oct 08, 2020
<u>RIOCIGUAT - ADEMPAS</u>						
N 204819	004	7173037	Dec 04, 2026	DS DP	ODE-53	Oct 08, 2020
<u>RIOCIGUAT - ADEMPAS</u>						
N 204819	005	7173037	Dec 04, 2026	DS DP	ODE-53	Oct 08, 2020
<u>RISEDRONATE SODIUM - ACTONEL</u>						
N 020835	005	7192938	May 06, 2023		U-353	
		7718634	May 06, 2023		U-662	
<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 - RISPERDAL CONSTA</u>						
N 021346	001	6667061	May 25, 2020	DP		

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<u>RISPERIDONE - RISPERDAL CONSTA</u>						
N 021346	002 6667061	May 25, 2020	DP			
<u>RISPERIDONE - RISPERDAL CONSTA</u>						
N 021346	003 6667061	May 25, 2020	DP			
<u>RISPERIDONE - PERSERIS KIT</u>						
N 210655	001 10010612	Feb 13, 2028	DP		NP	Jul 27, 2021
	10058554	Sep 26, 2026	U-2363			
	10376590	Feb 13, 2028	U-2608			
	10406160	Jun 26, 2026	DP U-2608			
	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		NP	Jul 27, 2021
	10058554	Sep 26, 2026	U-2363			
	10376590	Feb 13, 2028	U-2608			
	10406160	Jun 26, 2026	DP U-2608			
	9180197	Feb 13, 2028	DP			
	9186413	Feb 13, 2028	U-543			
	9597402	Sep 26, 2026	DP			
<u>RITONAVIR - NORVIR</u>						
N 020945	001 7141593	May 22, 2020	DP			
	7432294	May 22, 2020	DP			
<u>RITONAVIR - NORVIR</u>						
N 022417	001 7364752	Nov 10, 2020	DP U-688			
	8268349	Aug 25, 2024	DP			
	8399015	Aug 25, 2024	DP			
	8399015*PED	Feb 25, 2025				
	8470347	Sep 17, 2026	DP			
	8470347*PED	Mar 17, 2027				
	8691878	Aug 25, 2024	U-688			
	8691878*PED	Feb 25, 2025				
<u>RITONAVIR - NORVIR</u>						
N 209512	001				ODE-184	Jun 07, 2024
<u>RIVAROXABAN - XARELTO</u>						
N 022406	001 7157456	Aug 28, 2024	DS DP U-1301		D-168	Oct 27, 2020
	7157456	Aug 28, 2024	DS DP U-1302		I-810	Oct 11, 2022
	7585860	Dec 11, 2020	DS			
	7592339	Dec 11, 2020	U-1167			
	7592339	Dec 11, 2020	U-1200			
	7592339	Dec 11, 2020	U-1301			
	7592339	Dec 11, 2020	U-1302			
	7592339	Dec 11, 2020	U-1303			
	7592339	Dec 11, 2020	U-2142			
	7592339	Dec 11, 2020	U-2640			
	9415053	Nov 13, 2024	DP U-1167			
	9415053	Nov 13, 2024	DP U-1200			
	9415053	Nov 13, 2024	DP U-1301			
	9415053	Nov 13, 2024	DP U-1302			
	9415053	Nov 13, 2024	DP U-2142			
	9415053	Nov 13, 2024	DP U-2640			
	9539218	Feb 17, 2034	U-1953			
	9539218	Feb 17, 2034	U-1954			
	9539218	Feb 17, 2034	U-1955			
	9539218	Feb 17, 2034	U-1957			
	9539218	Feb 17, 2034	U-2143			
	9539218	Feb 17, 2034	U-2641			
<u>RIVAROXABAN - XARELTO</u>						
N 022406	002 7157456	Aug 28, 2024	DS DP U-1301		I-810	Oct 11, 2022
	7157456	Aug 28, 2024	DS DP U-1302			
	7585860	Dec 11, 2020	DS			
	7592339	Dec 11, 2020	U-1167			
	7592339	Dec 11, 2020	U-1200			

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<u>RIVAROXABAN - XARELTO</u>						
N 022406 002	7592339	Dec 11, 2020	U-1301			
	7592339	Dec 11, 2020	U-1302			
	7592339	Dec 11, 2020	U-1303			
	9415053	Nov 13, 2024	DP U-1167			
	9415053	Nov 13, 2024	DP U-1200			
	9415053	Nov 13, 2024	DP U-1301			
	9415053	Nov 13, 2024	DP U-1302			
	9415053	Nov 13, 2024	DP U-1303			
	9539218	Feb 17, 2034	U-1953			
	9539218	Feb 17, 2034	U-1954			
	9539218	Feb 17, 2034	U-1955			
	9539218	Feb 17, 2034	U-1956			
	9539218	Feb 17, 2034	U-1957			
<u>RIVAROXABAN - XARELTO</u>						
N 022406 003	7157456	Aug 28, 2024	DS DP U-1301		I-810	Oct 11, 2022
	7157456	Aug 28, 2024	DS DP U-1302			
	7585860	Dec 11, 2020	DS			
	7592339	Dec 11, 2020	U-1167			
	7592339	Dec 11, 2020	U-1200			
	7592339	Dec 11, 2020	U-1301			
	7592339	Dec 11, 2020	U-1302			
	7592339	Dec 11, 2020	U-1303			
	9415053	Nov 13, 2024	DP U-1167			
	9415053	Nov 13, 2024	DP U-1200			
	9415053	Nov 13, 2024	DP U-1301			
	9415053	Nov 13, 2024	DP U-1302			
	9539218	Feb 17, 2034	U-1953			
	9539218	Feb 17, 2034	U-1954			
	9539218	Feb 17, 2034	U-1955			
	9539218	Feb 17, 2034	U-1957			
<u>RIVAROXABAN - XARELTO</u>						
N 022406 004	7157456	Aug 28, 2024	DS DP		I-810	Oct 11, 2022
	7585860	Dec 11, 2020	DS		I-824	Oct 11, 2021
	7592339	Dec 11, 2020	U-2435			
	9415053	Nov 13, 2024	DP U-2435			
<u>ROFLUMILAST - DALIRESP</u>						
N 022522 001	5712298	Jan 27, 2020	DS DP U-1115		D-171	Jan 23, 2021
	8431154	Feb 19, 2023	DP		M-208	Aug 31, 2020
	8536206	Mar 08, 2024	U-1115			
	8604064	Mar 08, 2024	U-1115			
	8618142	Mar 08, 2024	DP			
	9468598	Feb 19, 2023	DP			
<u>ROFLUMILAST - DALIRESP</u>						
N 022522 002	5712298	Jan 27, 2020	DS DP U-1115		NS	Jan 23, 2021
	8431154	Feb 19, 2023	DP			
	8536206	Mar 08, 2024	U-1115			
	8604064	Mar 08, 2024	U-1115			
	8618142	Mar 08, 2024	DP			
	9468598	Feb 19, 2023	DP			
<u>ROLAPITANT HYDROCHLORIDE - VARUBI</u>						
N 206500 001	7049320	Dec 08, 2023	DS DP U-1741		NCE	Sep 01, 2020
	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			
	8796299	Dec 17, 2022	U-1741			
<u>ROLAPITANT HYDROCHLORIDE - VARUBI</u>						
N 208399 001	7049320	Dec 08, 2023	DS DP U-1741		NCE	Sep 01, 2020
	7981905	Apr 04, 2027	U-1741			
	8178550	Apr 04, 2027	DS DP			
	8404702	Apr 04, 2027	U-1741			
	8470842	Jan 18, 2029	U-1741			

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<u>ROLAPITANT HYDROCHLORIDE - VARUBI</u>						
N 208399 001	8796299	Dec 17, 2022	U-1741			
	9101615	Jul 14, 2032	U-1741			
<u>ROMIDEPSIN - ISTODAX</u>						
N 022393 001	7608280	Aug 22, 2021	DS			
	7611724	Aug 22, 2021	DS			
<u>ROPINIROLE HYDROCHLORIDE - REQUIP XL</u>						
N 022008 001	7927624	Dec 02, 2021	DP U-20		M-203	Mar 23, 2020
	8303986	Apr 12, 2021	DP			
<u>ROPINIROLE HYDROCHLORIDE - REQUIP XL</u>						
N 022008 002	7927624	Dec 02, 2021	DP U-20		M-203	Mar 23, 2020
	8303986	Apr 12, 2021	DP			
<u>ROPINIROLE HYDROCHLORIDE - REQUIP XL</u>						
N 022008 003	7927624	Dec 02, 2021	DP U-20		M-203	Mar 23, 2020
	8303986	Apr 12, 2021	DP			
<u>ROPINIROLE HYDROCHLORIDE - REQUIP XL</u>						
N 022008 004	7927624	Dec 02, 2021	DP U-20		M-203	Mar 23, 2020
	8303986	Apr 12, 2021	DP			
<u>ROPINIROLE HYDROCHLORIDE - REQUIP XL</u>						
N 022008 005	7927624	Dec 02, 2021	DP U-20		M-203	Mar 23, 2020
	8303986	Apr 12, 2021	DP			
<u>ROPINIROLE HYDROCHLORIDE - REQUIP XL</u>						
N 022008 006	7927624	Dec 02, 2021	DP U-20		M-203	Mar 23, 2020
	8303986	Apr 12, 2021	DP			
<u>ROPIVACAINE HYDROCHLORIDE - NAROPIN</u>						
N 020533 006	7828787	Oct 18, 2025	DP			
	7857802	Nov 28, 2026	DP			
	8118802	May 18, 2023	DP			
	8162915	May 23, 2024	DP			
<u>ROPIVACAINE HYDROCHLORIDE - NAROPIN</u>						
N 020533 007	7828787	Oct 18, 2025	DP			
	7857802	Nov 28, 2026	DP			
	8118802	May 18, 2023	DP			
	8162915	May 23, 2024	DP			
<u>ROSIGLITAZONE MALEATE - AVANDIA</u>						
N 021071 002	7358366	Apr 19, 2020	DS		Y	
	7358366*PED	Oct 19, 2020				
<u>ROSIGLITAZONE MALEATE - AVANDIA</u>						
N 021071 003	7358366	Apr 19, 2020	DS		Y	
	7358366*PED	Oct 19, 2020				
<u>ROSIGLITAZONE MALEATE - AVANDIA</u>						
N 021071 004	7358366	Apr 19, 2020	DS		Y	
	7358366*PED	Oct 19, 2020				
<u>ROSUVASTATIN CALCIUM - CRESTOR</u>						
N 021366 002	6316460	Aug 04, 2020	DP		ODE-118	May 27, 2023
	6858618	Dec 17, 2021	U-1032			
	6858618	Dec 17, 2021	U-1807			
	6858618	Dec 17, 2021	U-618			
	6858618*PED	Jun 17, 2022				
<u>ROSUVASTATIN CALCIUM - CRESTOR</u>						
N 021366 003	6316460	Aug 04, 2020	DP		ODE-118	May 27, 2023
	6858618	Dec 17, 2021	U-1032			
	6858618	Dec 17, 2021	U-1807			
	6858618	Dec 17, 2021	U-618			
	6858618*PED	Jun 17, 2022				

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<u>ROSUVASTATIN CALCIUM - CRESTOR</u>						
N 021366 004	6316460	Aug 04, 2020	DP		ODE-118	May 27, 2023
	6858618	Dec 17, 2021	U-1032			
	6858618	Dec 17, 2021	U-1807			
	6858618	Dec 17, 2021	U-618			
	6858618*PED	Jun 17, 2022				
<u>ROSUVASTATIN CALCIUM - CRESTOR</u>						
N 021366 005	6316460	Aug 04, 2020	DP		ODE-118	May 27, 2023
	6858618	Dec 17, 2021	U-618			
<u>ROSUVASTATIN CALCIUM - EZALLOR</u>						
N 208647 001	10413543	Feb 12, 2036	DP			
<u>ROSUVASTATIN CALCIUM - EZALLOR</u>						
N 208647 002	10413543	Feb 12, 2036	DP			
<u>ROSUVASTATIN CALCIUM - EZALLOR</u>						
N 208647 003	10413543	Feb 12, 2036	DP			
<u>ROSUVASTATIN CALCIUM - EZALLOR</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			
	6699498	Nov 27, 2020	DP			
	6884434	Mar 30, 2021	DP			
	8246979	Sep 01, 2027	DP U-1272			
	8246979	Sep 01, 2027	DP U-1273			
	8246980	Nov 27, 2025	DP			
	8617591	Jul 22, 2023	DP U-1474			
	9925150	Mar 01, 2032	DP			
<u>ROTIGOTINE - NEUPRO</u>						
N 021829 002	10130589	Dec 22, 2030	DP			
	10350174	Dec 22, 2030	DP			
	6699498	Nov 27, 2020	DP			
	6884434	Mar 30, 2021	DP			
	8246979	Sep 01, 2027	DP U-1272			
	8246979	Sep 01, 2027	DP U-1273			
	8246980	Nov 27, 2025	DP			
	8617591	Jul 22, 2023	DP U-1474			
	9925150	Mar 01, 2032	DP			
<u>ROTIGOTINE - NEUPRO</u>						
N 021829 003	10130589	Dec 22, 2030	DP			
	10350174	Dec 22, 2030	DP			
	6699498	Nov 27, 2020	DP			
	6884434	Mar 30, 2021	DP			
	8246979	Sep 01, 2027	DP U-1272			
	8246979	Sep 01, 2027	DP U-1273			
	8246980	Nov 27, 2025	DP			
	8617591	Jul 22, 2023	DP U-1474			
	9925150	Mar 01, 2032	DP			
<u>ROTIGOTINE - NEUPRO</u>						
N 021829 004	10130589	Dec 22, 2030	DP			
	10350174	Dec 22, 2030	DP			
	6699498	Nov 27, 2020	DP			
	6884434	Mar 30, 2021	DP			
	8246979	Sep 01, 2027	DP U-1272			
	8246979	Sep 01, 2027	DP U-1273			
	8246980	Nov 27, 2025	DP			
	8617591	Jul 22, 2023	DP U-1474			
	9925150	Mar 01, 2032	DP			
<u>ROTIGOTINE - NEUPRO</u>						
N 021829 005	10130589	Dec 22, 2030	DP			
	10350174	Dec 22, 2030	DP			
	6699498	Nov 27, 2020	DP			

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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	005 6884434	Mar 30, 2021	DP			
	8246979	Sep 01, 2027	DP U-1272			
	8246979	Sep 01, 2027	DP U-1273			
	8246980	Nov 27, 2025	DP			
	8617591	Jul 22, 2023	DP U-1474			
	9925150	Mar 01, 2032	DP			
<u>ROTIGOTINE - NEUPRO</u>						
N 021829	006 10130589	Dec 22, 2030	DP			
	10350174	Dec 22, 2030	DP			
	6699498	Nov 27, 2020	DP			
	6884434	Mar 30, 2021	DP			
	8246979	Sep 01, 2027	DP U-1272			
	8246979	Sep 01, 2027	DP U-1273			
	8246980	Nov 27, 2025	DP			
	8617591	Jul 22, 2023	DP U-1474			
	9925150	Mar 01, 2032	DP			
<u>RUCAPARIB CAMSYLATE - RUBRACA</u>						
N 209115	001 10130636	Aug 17, 2035	U-2012		I-772	Apr 06, 2021
	10130636	Aug 17, 2035	U-2101		NCE	Dec 19, 2021
	10130636	Aug 17, 2035	U-2273		ODE-126	Dec 19, 2023
	10278974	Feb 10, 2031	DP		ODE-168	Apr 06, 2025
	6495541	Jan 10, 2021	DS DP			
	7351701	Jul 23, 2024	U-2012			
	7351701	Jul 23, 2024	U-2101			
	7351701	Jul 23, 2024	U-2273			
	7531530	Jul 23, 2024	U-2012			
	7531530	Jul 23, 2024	U-2101			
	7531530	Jul 23, 2024	U-2273			
	8071579	Aug 12, 2027	U-2012			
	8071579	Aug 12, 2027	U-2101			
	8071579	Aug 12, 2027	U-2273			
	8143241	Aug 12, 2027	U-2012			
	8143241	Aug 12, 2027	U-2101			
	8143241	Aug 12, 2027	U-2273			
	8754072	Feb 10, 2031	DS DP			
	8859562	Aug 04, 2031	U-2012			
	8859562	Aug 04, 2031	U-2101			
	8859562	Aug 04, 2031	U-2273			
	9045487	Feb 10, 2031	DS DP			
	9861638	Feb 10, 2031	U-2012			
	9861638	Feb 10, 2031	U-2101			
	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		I-772	Apr 06, 2021
	10130636	Aug 17, 2035	U-2101		NCE	Dec 19, 2021
	10130636	Aug 17, 2035	U-2273		ODE-126	Dec 19, 2023
	10278974	Feb 10, 2031	DP		ODE-168	Apr 06, 2025
	6495541	Jan 10, 2021	DS DP			
	7351701	Jul 23, 2024	U-2012			
	7351701	Jul 23, 2024	U-2101			
	7351701	Jul 23, 2024	U-2273			
	7531530	Jul 23, 2024	U-2012			
	7531530	Jul 23, 2024	U-2101			
	7531530	Jul 23, 2024	U-2273			
	8071579	Aug 12, 2027	U-2012			
	8071579	Aug 12, 2027	U-2101			
	8071579	Aug 12, 2027	U-2273			
	8143241	Aug 12, 2027	U-2012			
	8143241	Aug 12, 2027	U-2101			
	8143241	Aug 12, 2027	U-2273			
	8754072	Feb 10, 2031	DS DP			
	8859562	Aug 04, 2031	U-2012			
	8859562	Aug 04, 2031	U-2101			
	8859562	Aug 04, 2031	U-2273			
	9045487	Feb 10, 2031	DS DP			
	9861638	Feb 10, 2031	U-2012			

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<u>RUCAPARIB CAMSYLATE - RUBRACA</u>						
N 209115 002	9861638	Feb 10, 2031	U-2101			
	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		I-772	Apr 06, 2021
	10130636	Aug 17, 2035	U-2101		NCE	Dec 19, 2021
	10130636	Aug 17, 2035	U-2273		ODE-126	Dec 19, 2023
	10278974	Feb 10, 2031	DP		ODE-168	Apr 06, 2025
	6495541	Jan 10, 2021	DS DP			
	7351701	Jul 23, 2024	U-2012			
	7351701	Jul 23, 2024	U-2101			
	7351701	Jul 23, 2024	U-2273			
	7531530	Jul 23, 2024	U-2012			
	7531530	Jul 23, 2024	U-2101			
	7531530	Jul 23, 2024	U-2273			
	8071579	Aug 12, 2027	U-2012			
	8071579	Aug 12, 2027	U-2101			
	8071579	Aug 12, 2027	U-2273			
	8143241	Aug 12, 2027	U-2012			
	8143241	Aug 12, 2027	U-2101			
	8143241	Aug 12, 2027	U-2273			
	8754072	Feb 10, 2031	DS DP			
	8859562	Aug 04, 2031	U-2012			
	8859562	Aug 04, 2031	U-2101			
	8859562	Aug 04, 2031	U-2273			
	9045487	Feb 10, 2031	DS DP			
	9861638	Feb 10, 2031	U-2012			
	9861638	Feb 10, 2031	U-2101			
	9861638	Feb 10, 2031	U-2273			
	9987285	Aug 17, 2035	DP			
<u>RUFINAMIDE - BANZEL</u>						
N 021911 001	6740669	Nov 14, 2022	DS DP			
	6740669*PED	May 14, 2023				
<u>RUFINAMIDE - BANZEL</u>						
N 021911 002	6740669	Nov 14, 2022	DS DP			
	6740669*PED	May 14, 2023				
<u>RUFINAMIDE - BANZEL</u>						
N 021911 003	6740669	Nov 14, 2022	DS DP			
	6740669*PED	May 14, 2023				
<u>RUFINAMIDE - BANZEL</u>						
N 201367 001	6740669	Nov 14, 2022	DS DP			
	6740669*PED	May 14, 2023				
<u>RUXOLITINIB PHOSPHATE - JAKAFI</u>						
N 202192 001	10016429	Jun 12, 2028	U-2536		I-799	May 24, 2022
	7598257	Dec 24, 2027	DS DP U-1201		ODE-238	May 24, 2026
	7598257	Dec 24, 2027	DS DP U-1622		ODE-79	Dec 04, 2021
	8415362	Dec 24, 2027	DS DP			
	8722693	Jun 12, 2028	DS DP			
	8822481	Jun 12, 2028	U-1573			
	8829013	Jun 12, 2028	U-1201			
	8829013	Jun 12, 2028	U-1622			
	9079912	Dec 12, 2026	U-1573			
	9079912	Dec 12, 2026	U-1721			
	9814722	Dec 12, 2026	U-2536			
<u>RUXOLITINIB PHOSPHATE - JAKAFI</u>						
N 202192 002	10016429	Jun 12, 2028	U-2536		I-799	May 24, 2022
	7598257	Dec 24, 2027	DS DP U-1201		ODE-238	May 24, 2026
	7598257	Dec 24, 2027	DS DP U-1622		ODE-79	Dec 04, 2021
	8415362	Dec 24, 2027	DS DP			
	8722693	Jun 12, 2028	DS DP			
	8822481	Jun 12, 2028	U-1573			
	8829013	Jun 12, 2028	U-1201			
	8829013	Jun 12, 2028	U-1622			

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<u>RUXOLITINIB PHOSPHATE - JAKAFI</u>						
N 202192 002	9079912	Dec 12, 2026	U-1573			
	9079912	Dec 12, 2026	U-1721			
	9814722	Dec 12, 2026	U-2536			
<u>RUXOLITINIB PHOSPHATE - JAKAFI</u>						
N 202192 003	10016429	Jun 12, 2028	U-2536		I-799	May 24, 2022
	7598257	Dec 24, 2027	DS DP U-1201		ODE-238	May 24, 2026
	7598257	Dec 24, 2027	DS DP U-1622		ODE-79	Dec 04, 2021
	8415362	Dec 24, 2027	DS DP			
	8722693	Jun 12, 2028	DS DP			
	8822481	Jun 12, 2028	U-1573			
	8829013	Jun 12, 2028	U-1201			
	8829013	Jun 12, 2028	U-1622			
	9079912	Dec 12, 2026	U-1573			
	9079912	Dec 12, 2026	U-1721			
	9814722	Dec 12, 2026	U-2536			
<u>RUXOLITINIB PHOSPHATE - JAKAFI</u>						
N 202192 004	10016429	Jun 12, 2028	U-2536		I-799	May 24, 2022
	7598257	Dec 24, 2027	DS DP U-1201		ODE-238	May 24, 2026
	7598257	Dec 24, 2027	DS DP U-1622		ODE-79	Dec 04, 2021
	8415362	Dec 24, 2027	DS DP			
	8722693	Jun 12, 2028	DS DP			
	8822481	Jun 12, 2028	U-1573			
	8829013	Jun 12, 2028	U-1201			
	8829013	Jun 12, 2028	U-1622			
	9079912	Dec 12, 2026	U-1573			
	9079912	Dec 12, 2026	U-1721			
	9814722	Dec 12, 2026	U-2536			
<u>RUXOLITINIB PHOSPHATE - JAKAFI</u>						
N 202192 005	10016429	Jun 12, 2028	U-2536		I-799	May 24, 2022
	7598257	Dec 24, 2027	DS DP U-1201		ODE-238	May 24, 2026
	7598257	Dec 24, 2027	DS DP U-1622		ODE-79	Dec 04, 2021
	8415362	Dec 24, 2027	DS DP			
	8722693	Jun 12, 2028	DS DP			
	8822481	Jun 12, 2028	U-1573			
	8829013	Jun 12, 2028	U-1201			
	8829013	Jun 12, 2028	U-1622			
	9079912	Dec 12, 2026	U-1573			
	9079912	Dec 12, 2026	U-1721			
	9814722	Dec 12, 2026	U-2536			
<u>SACROSIDASE - SUCRAID</u>						
N 020772 001	9255261	Feb 07, 2034	DS DP			
	9469847	Feb 07, 2034	DS DP			
	9849161	Feb 07, 2034	DS DP			
<u>SACUBITRIL; VALSARTAN - ENTRESTO</u>						
N 207620 001	7468390	Nov 27, 2023	DP		NCE	Jul 07, 2020
	7468390*PED	May 27, 2024			NPP	Oct 01, 2022
	8101659	Jan 14, 2023	DP		PED	Jan 07, 2021
	8101659*PED	Jul 14, 2023			PED	Apr 01, 2023
	8404744	Jan 14, 2023	DP			
	8404744*PED	Jul 14, 2023				
	8796331	Jan 14, 2023	U-1723			
	8796331*PED	Jul 14, 2023				
	8877938	May 27, 2027	DS DP			
	8877938*PED	Nov 27, 2027				
	9388134	Nov 08, 2026	U-1723			
	9388134*PED	May 08, 2027				
<u>SACUBITRIL; VALSARTAN - ENTRESTO</u>						
N 207620 002	7468390	Nov 27, 2023	DP		NCE	Jul 07, 2020
	7468390*PED	May 27, 2024			NPP	Oct 01, 2022
	8101659	Jan 14, 2023	DP		PED	Jan 07, 2021
	8101659*PED	Jul 14, 2023			PED	Apr 01, 2023
	8404744	Jan 14, 2023	DP			
	8404744*PED	Jul 14, 2023				
	8796331	Jan 14, 2023	U-1723			

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<u>SACUBITRIL; VALSARTAN - ENTRESTO</u>						
N 207620 002	8796331*PED	Jul 14, 2023				
	8877938	May 27, 2027	DS DP			
	8877938*PED	Nov 27, 2027				
	9388134	Nov 08, 2026			U-1723	
	9388134*PED	May 08, 2027				
<u>SACUBITRIL; VALSARTAN - ENTRESTO</u>						
N 207620 003	7468390	Nov 27, 2023		DP	NCE	Jul 07, 2020
	7468390*PED	May 27, 2024			NPP	Oct 01, 2022
	8101659	Jan 14, 2023		DP	PED	Jan 07, 2021
	8101659*PED	Jul 14, 2023			PED	Apr 01, 2023
	8404744	Jan 14, 2023		DP		
	8404744*PED	Jul 14, 2023				
	8796331	Jan 14, 2023			U-1723	
	8796331*PED	Jul 14, 2023				
	8877938	May 27, 2027		DS DP		
	8877938*PED	Nov 27, 2027				
	9388134	Nov 08, 2026			U-1723	
	9388134*PED	May 08, 2027				
<u>SAFINAMIDE MESYLATE - XADAGO</u>						
N 207145 001	8076515	Dec 10, 2028	DS DP	U-1993	NCE	Mar 21, 2022
	8278485	Jun 08, 2027	DS	U-1993		
	8283380	Sep 01, 2027		U-1993		
<u>SAFINAMIDE MESYLATE - XADAGO</u>						
N 207145 002	8076515	Dec 10, 2028	DS DP	U-1993	NCE	Mar 21, 2022
	8278485	Jun 08, 2027	DS	U-1993		
	8283380	Sep 01, 2027		U-1993		
<u>SAPROPTERIN DIHYDROCHLORIDE - KUVAN</u>						
N 022181 001	7566462	Nov 16, 2025		DP		
	7566462*PED	May 16, 2026				
	7566714	Nov 17, 2024			U-989	
	7566714*PED	May 17, 2025				
	7612073	Nov 17, 2024			U-1010	
	7612073*PED	May 17, 2025				
	7727987	Nov 17, 2024		DP		
	7727987*PED	May 17, 2025				
	7947681	Nov 17, 2024			U-1156	
	7947681*PED	May 17, 2025				Y
	8003126	Nov 16, 2025				
	8003126*PED	May 16, 2026				
	8067416	Nov 17, 2024			U-989	
	8067416*PED	May 17, 2025				
	8318745	Nov 17, 2024		DP		
	8318745*PED	May 17, 2025				
	9433624	Nov 17, 2024			U-1589	
	RE43797	Nov 17, 2024			U-1156	
	RE43797*PED	May 17, 2025				
<u>SAPROPTERIN DIHYDROCHLORIDE - KUVAN</u>						
N 205065 001	7566714	Nov 17, 2024			U-1589	
	7566714*PED	May 17, 2025				
	7612073	Nov 17, 2024			U-1010	
	7612073*PED	May 17, 2025				
	8067416	Nov 17, 2024			U-1589	
	8067416*PED	May 17, 2025				
	9216178	Nov 01, 2032		DP		
	9433624	Nov 17, 2024			U-1589	
	RE43797	Nov 17, 2024			U-1590	
	RE43797*PED	May 17, 2025				
<u>SAPROPTERIN DIHYDROCHLORIDE - KUVAN</u>						
N 205065 002	7566714	Nov 17, 2024			U-1589	
	7566714*PED	May 17, 2025				
	7612073	Nov 17, 2024			U-1010	
	7612073*PED	May 17, 2025				
	8067416	Nov 17, 2024			U-1589	
	8067416*PED	May 17, 2025				

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<u>SAPROPTERIN DIHYDROCHLORIDE - KUVAN</u>						
N 205065 002	9216178	Nov 01, 2032	DP			
	9433624	Nov 17, 2024	U-1589			
	RE43797	Nov 17, 2024	U-1590			
	RE43797*PED	May 17, 2025				
<u>SARECYCLINE HYDROCHLORIDE - SEYSARA</u>						
N 209521 001	8318706	May 01, 2031	DS DP U-2405		NCE	Oct 01, 2023
	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		NCE	Oct 01, 2023
	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		NCE	Oct 01, 2023
	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		M-198	Feb 27, 2020
	RE44186	Jul 31, 2023	DS DP U-1837			
	RE44186	Jul 31, 2023	DS DP U-995			
<u>SAXAGLIPTIN HYDROCHLORIDE - ONGLYZA</u>						
N 022350 002	7951400	Nov 30, 2028	DP		M-198	Feb 27, 2020
	RE44186	Jul 31, 2023	DS DP U-1837			
	RE44186	Jul 31, 2023	DS DP U-995			
<u>SECNIDAZOLE - SOLOSEC</u>						
N 209363 001	10335390	Sep 04, 2035	U-2583		NCE GAIN	Sep 15, 2022 Sep 15, 2027
<u>SELENIOS ACID - SELENIOS ACID</u>						
N 209379 001					NCE	Apr 30, 2024
<u>SELEXIPAG - UPTRAVI</u>						
N 207947 001	7205302	Apr 04, 2023	DS DP U-1797		NCE	Dec 21, 2020
	8791122	Aug 01, 2030	DS DP		ODE-106	Dec 21, 2022
	9173881	Aug 12, 2029	U-1798			
	9284280	Jun 25, 2030	U-1831			
<u>SELEXIPAG - UPTRAVI</u>						
N 207947 002	7205302	Apr 04, 2023	DS DP U-1797		NCE	Dec 21, 2020
	8791122	Aug 01, 2030	DS DP		ODE-106	Dec 21, 2022
	9173881	Aug 12, 2029	U-1798			
	9284280	Jun 25, 2030	U-1831			
<u>SELEXIPAG - UPTRAVI</u>						
N 207947 003	7205302	Apr 04, 2023	DS DP U-1797		NCE	Dec 21, 2020
	8791122	Aug 01, 2030	DS DP		ODE-106	Dec 21, 2022
	9173881	Aug 12, 2029	U-1798			
	9284280	Jun 25, 2030	U-1831			
<u>SELEXIPAG - UPTRAVI</u>						
N 207947 004	7205302	Apr 04, 2023	DS DP U-1797		NCE	Dec 21, 2020
	8791122	Aug 01, 2030	DS DP		ODE-106	Dec 21, 2022
	9173881	Aug 12, 2029	U-1798			
	9284280	Jun 25, 2030	U-1831			
<u>SELEXIPAG - UPTRAVI</u>						
N 207947 005	7205302	Apr 04, 2023	DS DP U-1797		NCE	Dec 21, 2020
	8791122	Aug 01, 2030	DS DP		ODE-106	Dec 21, 2022

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<u>SELEXIPAG - UPTRAVI</u>						
N 207947 005	9173881	Aug 12, 2029		U-1798		
	9284280	Jun 25, 2030		U-1831		
<u>SELEXIPAG - UPTRAVI</u>						
N 207947 006	7205302	Apr 04, 2023	DS DP	U-1797	NCE	Dec 21, 2020
	8791122	Aug 01, 2030	DS DP		ODE-106	Dec 21, 2022
	9173881	Aug 12, 2029		U-1798		
	9284280	Jun 25, 2030		U-1831		
<u>SELEXIPAG - UPTRAVI</u>						
N 207947 007	7205302	Apr 04, 2023	DS DP	U-1797	NCE	Dec 21, 2020
	8791122	Aug 01, 2030	DS DP		ODE-106	Dec 21, 2022
	9173881	Aug 12, 2029		U-1798		
	9284280	Jun 25, 2030		U-1831		
<u>SELEXIPAG - UPTRAVI</u>						
N 207947 008	7205302	Apr 04, 2023	DS DP	U-1797	NCE	Dec 21, 2020
	8791122	Aug 01, 2030	DS DP		ODE-106	Dec 21, 2022
	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	NCE	Jul 03, 2024
	10544108	Jul 26, 2032		U-2584	ODE-257	Jul 03, 2026
	8999996	Sep 15, 2032	DS DP			
	9079865	Jul 26, 2032		U-2584		
	9714226	Jul 26, 2032	DS DP			
<u>SEMAGLUTIDE - OZEMPIC</u>						
N 209637 001	10220155	Jul 17, 2026		DP	I-822	Jan 16, 2023
	10335462	Jun 21, 2033		U-2580	NCE	Dec 05, 2022
	10357616	Jan 20, 2026		DP		
	10376652	Jan 20, 2026		DP		
	6899699	Jan 02, 2022		DP		
	7762994	May 23, 2024		DP		
	8114833	Aug 13, 2025		DP		
	8129343	Jan 29, 2029	DS DP	U-2202		
	8536122	Mar 20, 2026	DS DP	U-2202		
	8579869	Jun 30, 2023		DP		
	8672898	Jan 02, 2022		DP		
	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		
	9486588	Jan 02, 2022		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					I-822	Jan 16, 2023
<u>SEMAGLUTIDE - RYBELSUS</u>						
N 213051 001	10086047	Dec 16, 2031		DP	M-252	Jan 16, 2023
	10278923	May 02, 2034		U-2628	NCE	Dec 05, 2022
	8129343	Jan 29, 2029	DS DP	U-2628	NP	Sep 20, 2022
	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	M-252	Jan 16, 2023
	10278923	May 02, 2034		U-2628	NCE	Dec 05, 2022
	8129343	Jan 29, 2029	DS DP	U-2628	NP	Sep 20, 2022
	8536122	Mar 20, 2026	DS DP	U-2628		
	9278123	Dec 16, 2031		DP	U-2628	

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<u>SEMAGLUTIDE - RYBELSUS</u>						
N 213051	003 10086047	Dec 16, 2031	DP		M-252	Jan 16, 2023
	10278923	May 02, 2034	U-2628		NCE	Dec 05, 2022
	8129343	Jan 29, 2029	DS DP U-2628		NP	Sep 20, 2022
	8536122	Mar 20, 2026	DS DP U-2628			
	9278123	Dec 16, 2031	DP U-2628			
<u>SERTRALINE HYDROCHLORIDE - ZOLOFT</u>						
N 020990	001 7067555*PED	Apr 11, 2020				
<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			
<u>SEVELAMER CARBONATE - RENVELA</u>						
N 022318	002 9095509	Dec 06, 2030	DP			
<u>SEVELAMER HYDROCHLORIDE - RENAGEL</u>						
N 021179	001 6733780	Oct 18, 2020	DP			
<u>SEVELAMER HYDROCHLORIDE - RENAGEL</u>						
N 021179	002 6733780	Oct 18, 2020	DP			
<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 6699871	Jul 26, 2022	DS DP U-1188			
	7125873	Jul 26, 2022	DP U-1189			
	7125873	Jul 26, 2022	DP U-1190			
	7125873	Jul 26, 2022	DP U-1192			
	7125873	Jul 26, 2022	DP U-1193			
	7326708	Apr 11, 2026	DS DP U-1188			
	8168637	Jun 26, 2022	DP U-1188			
<u>SIMVASTATIN; SITAGLIPTIN PHOSPHATE - JUVISYNC</u>						
N 202343	002 6699871	Jul 26, 2022	DS DP U-1188			
	7125873	Jul 26, 2022	DP U-1189			
	7125873	Jul 26, 2022	DP U-1190			
	7125873	Jul 26, 2022	DP U-1192			
	7125873	Jul 26, 2022	DP U-1193			
	7326708	Apr 11, 2026	DS DP U-1188			
	8168637	Jun 26, 2022	DP U-1188			
<u>SIMVASTATIN; SITAGLIPTIN PHOSPHATE - JUVISYNC</u>						
N 202343	003 6699871	Jul 26, 2022	DS DP U-1188			
	7125873	Jul 26, 2022	DP U-1189			
	7125873	Jul 26, 2022	DP U-1190			
	7125873	Jul 26, 2022	DP U-1192			
	7125873	Jul 26, 2022	DP U-1193			
	7326708	Apr 11, 2026	DS DP U-1188			
	8168637	Jun 26, 2022	DP U-1188			

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<u>SIMVASTATIN; SITAGLIPTIN PHOSPHATE - JUVISYNC</u>						
N 202343 004	6699871	Jul 26, 2022	DS DP U-1188			
	7125873	Jul 26, 2022	DP U-1189			
	7125873	Jul 26, 2022	DP U-1190			
	7125873	Jul 26, 2022	DP U-1192			
	7125873	Jul 26, 2022	DP U-1193			
	7326708	Apr 11, 2026	DS DP U-1188			
	8168637	Jun 26, 2022	DP U-1188			
<u>SIMVASTATIN; SITAGLIPTIN PHOSPHATE - JUVISYNC</u>						
N 202343 005	6699871	Jul 26, 2022	DS DP U-1188			
	7125873	Jul 26, 2022	DP U-1189			
	7125873	Jul 26, 2022	DP U-1190			
	7125873	Jul 26, 2022	DP U-1192			
	7125873	Jul 26, 2022	DP U-1193			
	7326708	Apr 11, 2026	DS DP U-1188			
	8168637	Jun 26, 2022	DP U-1188			
<u>SIMVASTATIN; SITAGLIPTIN PHOSPHATE - JUVISYNC</u>						
N 202343 006	6699871	Jul 26, 2022	DS DP U-1188			
	7125873	Jul 26, 2022	DP U-1189			
	7125873	Jul 26, 2022	DP U-1190			
	7125873	Jul 26, 2022	DP U-1192			
	7125873	Jul 26, 2022	DP U-1193			
	7326708	Apr 11, 2026	DS DP U-1188			
	8168637	Jun 26, 2022	DP U-1188			
<u>SINCALIDE - KINEVAC</u>						
N 017697 001	6803046	Aug 16, 2022	DP			
<u>SINECATECHINS - VEREGEN</u>						
N 021902 001	10434059	Nov 18, 2022	DP U-172			
	5795911	Oct 31, 2020	U-172			
	7858662	Oct 02, 2026	DP U-172			
	9770406	Jul 12, 2025	DP U-172			
<u>SIPONIMOD FUMARIC ACID - MAYZENT</u>						
N 209884 001	7939519	May 19, 2024	DS DP		NCE	Mar 26, 2024
	8492441	Nov 30, 2030	U-2511			
<u>SIPONIMOD FUMARIC ACID - MAYZENT</u>						
N 209884 002	7939519	May 19, 2024	DS DP		NCE	Mar 26, 2024
	8492441	Nov 30, 2030	U-2511			
<u>SIROLIMUS - RAPAMUNE</u>						
N 021083 001					ODE-92	May 28, 2022
<u>SIROLIMUS - RAPAMUNE</u>						
N 021110 001					ODE-92	May 28, 2022
<u>SIROLIMUS - RAPAMUNE</u>						
N 021110 002					ODE-92	May 28, 2022
<u>SIROLIMUS - RAPAMUNE</u>						
N 021110 003					ODE-92	May 28, 2022
<u>SIROLIMUS - RAPAMUNE</u>						
N 021110 004					ODE-92	May 28, 2022
<u>SITAGLIPTIN PHOSPHATE - JANUVIA</u>						
N 021995 001	6699871	Jul 26, 2022	DS DP U-774		M-244	Aug 12, 2022
	7125873	Jul 26, 2022	U-1036			
	7125873	Jul 26, 2022	U-1037			
	7125873	Jul 26, 2022	U-1038			
	7125873	Jul 26, 2022	U-775			
	7326708	Nov 24, 2026	DS DP U-802			
<u>SITAGLIPTIN PHOSPHATE - JANUVIA</u>						
N 021995 002	6699871	Jul 26, 2022	DS DP U-774		M-244	Aug 12, 2022
	7125873	Jul 26, 2022	U-1036			
	7125873	Jul 26, 2022	U-1037			
	7125873	Jul 26, 2022	U-1038			

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<u>SITAGLIPTIN PHOSPHATE - JANUVIA</u>						
N 021995 002	7125873	Jul 26, 2022		U-775		
	7326708	Nov 24, 2026	DS DP	U-802		
<u>SITAGLIPTIN PHOSPHATE - JANUVIA</u>						
N 021995 003	6699871	Jul 26, 2022	DS DP	U-774	M-244	Aug 12, 2022
	7125873	Jul 26, 2022		U-1036		
	7125873	Jul 26, 2022		U-1037		
	7125873	Jul 26, 2022		U-1038		
	7125873	Jul 26, 2022		U-775		
	7326708	Nov 24, 2026	DS DP	U-802		
<u>SODIUM NITRITE - SODIUM NITRITE</u>						
N 203922 001	8568793	Dec 24, 2031	DS DP			
<u>SODIUM NITRITE; SODIUM THIOSULFATE - NITHIODOTE</u>						
N 201444 001	8496973	Mar 29, 2031	DS DP	U-1419		
	8568793	Dec 24, 2031	DS DP			
	9345724	Mar 29, 2031	DS DP	U-2015		
	9585912	Mar 29, 2031	DS DP			
<u>SODIUM OXYBATE - XYREM</u>						
N 021196 001	10213400	Mar 15, 2033		U-2499	NPP	Oct 26, 2021
	6780889	Jul 04, 2020	DP		ODE-231	Oct 26, 2025
	6780889*PED	Jan 04, 2021			PED	Apr 26, 2022
	7262219	Jul 04, 2020	DP			
	7262219*PED	Jan 04, 2021				
	7668730	Jun 16, 2024		U-1110	Y	
	7668730*PED	Dec 16, 2024				
	7851506*PED	Jun 22, 2020				
	8263650*PED	Jun 22, 2020				
	8324275*PED	Jun 22, 2020				
	8731963	Dec 17, 2022		U-1110		
	8731963*PED	Jun 17, 2023				
	8772306	Mar 15, 2033		U-1532		
	8772306*PED	Sep 15, 2033				
	8859619*PED	Jun 22, 2020				
	8952062*PED	Jun 22, 2020				
	9050302	Mar 15, 2033		U-1532		
	9050302*PED	Sep 15, 2033				
	9486426	Mar 15, 2033		U-1532		
	9486426*PED	Sep 15, 2033				
	9539330*PED	Jun 22, 2020				
<u>SODIUM PHOSPHATE, DIBASIC, ANHYDROUS; SODIUM PHOSPHATE, MONOBASIC, MONOHYDRATE - OSMOPREP</u>						
N 021892 001	7687075	Jun 22, 2028	DS DP			
<u>SODIUM THIOSULFATE - SODIUM THIOSULFATE</u>						
N 203923 001	8496973	Mar 29, 2031	DS DP	U-1419		
	9345724	Mar 29, 2031	DS DP	U-2015		
	9585912	Mar 29, 2031	DS DP			
<u>SODIUM ZIRCONIUM CYCLOSILICATE - LOKELMA</u>						
N 207078 001	10300087	Oct 14, 2035	DS	U-2312	NCE	May 18, 2023
	10335432	Feb 10, 2032		U-2312		
	10398730	Feb 10, 2032		U-2312		
	10413569	Feb 10, 2032	DS			
	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	NCE	May 18, 2023
	10398730	Feb 10, 2032		U-2312		
	10413569	Feb 10, 2032	DS			
	8802152	Apr 19, 2032	DS			
	8808750	Feb 10, 2032		U-2312		

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<u>SODIUM ZIRCONIUM CYCLOSILICATE - LOKELMA</u>						
N 207078 002	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		NPP	Apr 07, 2020
	7964580*PED	Sep 26, 2029			ODE-135	Apr 07, 2024
	8334270	Mar 21, 2028	DS DP U-1470		PED	Oct 07, 2024
	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				
	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			
	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				
<u>SOFOSEBUVIR - 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			

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<u>SOFOSBUVIR - SOVALDI</u>						
N 212480 002	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	D-177	Nov 15, 2022
	7964580	Mar 26, 2029	DS DP U-1470		NCE	Jun 28, 2021
	7964580*PED	Sep 26, 2029			NPP	Aug 01, 2020
	8334270	Mar 21, 2028	DS DP U-1470			
	8334270*PED	Sep 21, 2028				
	8575135	Nov 16, 2032	DS DP U-1470			
	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		
	8889159	Mar 26, 2029	DP U-1470			
	8889159*PED	Sep 26, 2029				
	8921341	Nov 16, 2032	DS DP U-1470			
	8940718	Nov 16, 2032	DS DP U-1470			
	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			
<u>SOFOSBUVIR; VELPATASVIR; VOXILAPREVIR - VOSEVI</u>						
N 209195 001	7964580	Mar 26, 2029	DS DP U-2039		NCE	Jul 18, 2022
	7964580	Mar 26, 2029	DS DP U-2040			
	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 05, 2033	DS DP U-2039			
	8575135	Nov 05, 2033	DS DP U-2040			
	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			
	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			
	8940718	Nov 16, 2032	DS DP U-2039			
	8940718	Nov 16, 2032	DS DP 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			
	9585906	Mar 21, 2028	DS DP U-2039			

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<u>SOFOSBUVIR; VELPATASVIR; VOXILAPREVIR - VOSEVI</u>						
N 209195 001	9585906	Mar 21, 2028	DS DP U-2040			
	9868745	Nov 16, 2032	DS DP			
<u>SOLRIAMFETOL HYDROCHLORIDE - SUNOSI</u>						
N 211230 001	10195151	Sep 05, 2037	DP		NCE	Jun 17, 2024
	10351517	Jun 07, 2026	U-2548		ODE-254	Jun 17, 2026
	10512609	Sep 05, 2037	U-2548			
	8440715	Aug 25, 2027	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		NCE	Jun 17, 2024
	10351517	Jun 07, 2026	U-2548		ODE-254	Jun 17, 2026
	10512609	Sep 05, 2037	U-2548			
	8440715	Aug 25, 2027	U-2548			
	8877806	Jun 07, 2026	U-2548			
	9604917	Jun 07, 2026	U-2548			
<u>SOMATROPIN - NORDITROPIN NORDIFLEX</u>						
N 021148 004	RE41956	Jan 21, 2021	DP			
<u>SOMATROPIN - NORDITROPIN NORDIFLEX</u>						
N 021148 005	RE41956	Jan 21, 2021	DP			
<u>SOMATROPIN - NORDITROPIN NORDIFLEX</u>						
N 021148 006	RE41956	Jan 21, 2021	DP			
<u>SOMATROPIN - NORDITROPIN NORDIFLEX</u>						
N 021148 007	RE41956	Jan 21, 2021	DP			
<u>SOMATROPIN - NORDITROPIN FLEXPRO</u>						
N 021148 008	10220155	Jul 17, 2026	DP			
	10357616	Jan 20, 2026	DP			
	10376652	Jan 20, 2026	DP			
	6899699	Jan 02, 2022	DP			
	7762994	May 23, 2024	DP			
	8579869	Jun 30, 2023	DP			
	8672898	Jan 02, 2022	DP			
	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			
	9486588	Jan 02, 2022	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>SOMATROPIN - NORDITROPIN FLEXPRO</u>						
N 021148 009	10220155	Jul 17, 2026	DP			
	10357616	Jan 20, 2026	DP			
	10376652	Jan 20, 2026	DP			
	6899699	Jan 02, 2022	DP			
	7762994	May 23, 2024	DP			
	8579869	Jun 30, 2023	DP			
	8672898	Jan 02, 2022	DP			
	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			
	9486588	Jan 02, 2022	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			

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APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>SOMATROPIN - NORDITROPIN FLEXPRO</u>						
N 021148 009	10220155	Jul 17, 2026	DP			
	10357616	Jan 20, 2026	DP			
	10376652	Jan 20, 2026	DP			
	6899699	Jan 02, 2022	DP			
	7762994	May 23, 2024	DP			
	8579869	Jun 30, 2023	DP			
	8672898	Jan 02, 2022	DP			
	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			
	9486588	Jan 02, 2022	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>SOMATROPIN - NORDITROPIN FLEXPRO</u>						
N 021148 010	10220155	Jul 17, 2026	DP			
	10357616	Jan 20, 2026	DP			
	10376652	Jan 20, 2026	DP			
	6899699	Jan 02, 2022	DP			
	7762994	May 23, 2024	DP			
	8579869	Jun 30, 2023	DP			
	8672898	Jan 02, 2022	DP			
	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			
	9486588	Jan 02, 2022	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>SOMATROPIN - NORDITROPIN FLEXPRO</u>						
N 021148 011	10220155	Jul 17, 2026	DP			
	10357616	Jan 20, 2026	DP			
	10376652	Jan 20, 2026	DP			
	6899699	Jan 02, 2022	DP			
	7762994	May 23, 2024	DP			
	8579869	Jun 30, 2023	DP			
	8672898	Jan 02, 2022	DP			
	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			
	9486588	Jan 02, 2022	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>SONIDEGIB PHOSPHATE - ODOMZO</u>						
N 205266 001	8063043	Sep 15, 2029	DS DP		NCE	Jul 24, 2020
	8178563	Feb 06, 2029	DS	U-1722		
<u>SORAFENIB TOSYLATE - NEXAVAR</u>						
N 021923 001	7235576	Jan 12, 2020	DS DP		ODE-56	Nov 22, 2020
	7351834	Jan 12, 2020	DS			
	7897623	Jan 12, 2020	DP			
	8124630	Jan 12, 2020		U-1459		
	8618141	Feb 11, 2023		U-1480		
	8841330	Jan 12, 2020		U-1696		

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APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<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>SOTALOL HYDROCHLORIDE - SOTYLIZE</u>						
N 205108	001	10206895	Apr 01, 2034	DP U-2096		
		10206895	Apr 01, 2034	DP U-2494		
		9724297	Aug 31, 2035	DP U-2096		
<u>SPINOSAD - NATROBA</u>						
N 022408	001	6063771	Jul 25, 2023	DP U-1670		
		7030095	Jul 02, 2021	DP U-1105		
<u>SPIRONOLACTONE - CAROSPIR</u>						
N 209478	001	10493083	Oct 28, 2036	DP		
		9757394	Oct 28, 2036	DP U-2109		
<u>STAVUDINE - ZERIT XR</u>						
N 021453	001	7135465	Feb 18, 2023	DP U-167		
<u>STAVUDINE - ZERIT XR</u>						
N 021453	002	7135465	Feb 18, 2023	DP U-167		
<u>STAVUDINE - ZERIT XR</u>						
N 021453	003	7135465	Feb 18, 2023	DP U-167		
<u>STAVUDINE - ZERIT XR</u>						
N 021453	004	7135465	Feb 18, 2023	DP U-167		
<u>STIRIPENTOL - DIACOMIT</u>						
N 206709	001				NCE ODE-198	Aug 20, 2023 Aug 20, 2025
<u>STIRIPENTOL - DIACOMIT</u>						
N 206709	002				NCE ODE-198	Aug 20, 2023 Aug 20, 2025
<u>STIRIPENTOL - DIACOMIT</u>						
N 207223	001				NCE ODE-198	Aug 20, 2023 Aug 20, 2025
<u>STIRIPENTOL - DIACOMIT</u>						
N 207223	002				NCE ODE-198	Aug 20, 2023 Aug 20, 2025
<u>SUCROFERRIC OXYHYDROXIDE - VELPHORO</u>						
N 205109	001	6174442	Jun 12, 2020	DS U-1468		
		9561251	Jan 23, 2030	DP U-1468		
<u>SUFENTANIL CITRATE - DSUVIA</u>						
N 209128	001	10245228	Jan 05, 2027	DP U-1351	NP	Nov 02, 2021
		10342762	Jan 05, 2027	DP		
		10507180	Jan 05, 2027	DP U-1351		
		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		

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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	6949527	Jan 27, 2021	U-1795	NCE	Dec 15, 2020
		7265099	Aug 07, 2020	U-1795		
		RE44733	Jan 27, 2021	DS DP U-1794		
<u>SUGAMMADEX SODIUM - BRIDION</u>						
N 022225	002	6949527	Jan 27, 2021	U-1795	NCE	Dec 15, 2020
		7265099	Aug 07, 2020	U-1795		
		RE44733	Jan 27, 2021	DS DP U-1794		
<u>SULFUR HEXAFLUORIDE LIPID-TYPE A MICROSPHERES - LUMASON</u>						
N 203684	001	10232061	Jul 06, 2038	DP	NPP	Nov 13, 2022
		10335502	Jul 06, 2038	DP		
<u>SUMATRIPTAN - TOSYMRA</u>						
N 210884	001	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		
		8118771	Aug 10, 2023	DP		
		8241243	Aug 10, 2023	DP		
		8241244	Nov 21, 2022	DP		
		8267903	Mar 18, 2023	DP		
		8287489	Dec 06, 2024	DP		
		8343130	Oct 18, 2022	DP		
		8491524	Nov 21, 2022	DP		
<u>SUMATRIPTAN SUCCINATE - ALSUMA</u>						
N 022377	001	7811254	Aug 26, 2027	DP U-1083		
<u>SUMATRIPTAN SUCCINATE - ZECUITY</u>						
N 202278	001	6745071	Feb 21, 2023	DP		
		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		
		6715485	Mar 03, 2020	DP		
		7975690	Aug 18, 2025	DP U-1809		
		8047202	Jul 02, 2023	DP		
		8327844	Oct 03, 2023	U-1809		
		8550073	Oct 22, 2029	DP		
		8555877	Mar 03, 2020	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		
		9119932	Apr 23, 2024	DP		
		9649456	Oct 21, 2030	DP U-1719		

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<u>SUMATRIPTAN SUCCINATE - ONZETRA XSAIL</u>						
N 206099 001	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			
<u>SUNITINIB MALATE - SUTENT</u>						
N 021938 001	6573293	Feb 15, 2021	DS DP U-1154		I-755	Nov 16, 2020
	6573293	Feb 15, 2021	DS DP U-2171		PED	May 16, 2021
	6573293*PED	Aug 15, 2021				
	7125905	Feb 15, 2021	DS DP			
	7125905*PED	Aug 15, 2021				
	7211600	Dec 22, 2020	U-883			
	7211600*PED	Jun 22, 2021				
<u>SUNITINIB MALATE - SUTENT</u>						
N 021938 002	6573293	Feb 15, 2021	DS DP U-1154		I-755	Nov 16, 2020
	6573293	Feb 15, 2021	DS DP U-2171		PED	May 16, 2021
	6573293*PED	Aug 15, 2021				
	7125905	Feb 15, 2021	DS DP			
	7125905*PED	Aug 15, 2021				
	7211600	Dec 22, 2020	U-883			
	7211600*PED	Jun 22, 2021				
<u>SUNITINIB MALATE - SUTENT</u>						
N 021938 003	6573293	Feb 15, 2021	DS DP U-1154		I-755	Nov 16, 2020
	6573293	Feb 15, 2021	DS DP U-2171		PED	May 16, 2021
	6573293*PED	Aug 15, 2021				
	7125905	Feb 15, 2021	DS DP			
	7125905*PED	Aug 15, 2021				
	7211600	Dec 22, 2020	U-883			
	7211600*PED	Jun 22, 2021				
<u>SUNITINIB MALATE - SUTENT</u>						
N 021938 004	6573293	Feb 15, 2021	DS DP U-1154		I-755	Nov 16, 2020
	6573293	Feb 15, 2021	DS DP U-2171		PED	May 16, 2021
	6573293*PED	Aug 15, 2021				
	7125905	Feb 15, 2021	DS DP			
	7125905*PED	Aug 15, 2021				
	7211600	Dec 22, 2020	U-883			
	7211600*PED	Jun 22, 2021				
<u>SUVOREXANT - BELSOMRA</u>						
N 204569 001	10098892	May 29, 2033	DP		M-253	Jan 29, 2023
	7951797	Nov 20, 2029	DS DP U-620			
<u>SUVOREXANT - BELSOMRA</u>						
N 204569 002	10098892	May 29, 2033	DP		M-253	Jan 29, 2023
	7951797	Nov 20, 2029	DS DP U-620			
<u>SUVOREXANT - BELSOMRA</u>						
N 204569 003	10098892	May 29, 2033	DP		M-253	Jan 29, 2023
	7951797	Nov 20, 2029	DS DP U-620			
<u>SUVOREXANT - BELSOMRA</u>						
N 204569 004	10098892	May 29, 2033	DP		M-253	Jan 29, 2023
	7951797	Nov 20, 2029	DS DP U-620			
<u>TACROLIMUS - ENVARUSUS XR</u>						
N 206406 001	10166190	May 30, 2028	DP		ODE-94	Jul 10, 2022
	7994214	Aug 30, 2024	DP			
	8486993	Aug 30, 2024	DP U-1752			
	8586084	Aug 30, 2024	U-1752			
	8591946	Aug 30, 2024	DP			
	8617599	Aug 30, 2024	DP			
	8623410	Aug 30, 2024	DP			
	8623411	Aug 30, 2024	U-1752			
	8664239	Aug 30, 2028	U-1752			
	8664239	Aug 30, 2028	U-2677			
	8664239	Aug 30, 2028	U-2678			

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<u>TACROLIMUS - ENVARUSUS XR</u>						
N 206406 001	8685998	Aug 30, 2028	DP U-1752			
	8685998	Aug 30, 2028	DP U-2677			
	8685998	Aug 30, 2028	DP U-2678			
	8889185	Aug 30, 2024	U-1752			
	8889186	Aug 30, 2024	U-1752			
	9161907	Aug 30, 2024	DP U-1752			
	9549918	May 30, 2028	DP			
	9757362	Aug 30, 2024	DP			
	9763920	Aug 30, 2024	DP			
<u>TACROLIMUS - ENVARUSUS XR</u>						
N 206406 002	10166190	May 30, 2028	DP		ODE-94	Jul 10, 2022
	7994214	Aug 30, 2024	DP			
	8486993	Aug 30, 2024	DP U-1752			
	8586084	Aug 30, 2024	U-1752			
	8591946	Aug 30, 2024	DP			
	8617599	Aug 30, 2024	DP			
	8623410	Aug 30, 2024	DP			
	8623411	Aug 30, 2024	U-1752			
	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			
	8889185	Aug 30, 2024	U-1752			
	8889186	Aug 30, 2024	U-1752			
	9161907	Aug 30, 2024	DP U-1752			
	9549918	May 30, 2028	DP			
	9757362	Aug 30, 2024	DP			
	9763920	Aug 30, 2024	DP			
<u>TACROLIMUS - ENVARUSUS XR</u>						
N 206406 003	10166190	May 30, 2028	DP		ODE-94	Jul 10, 2022
	7994214	Aug 30, 2024	DP			
	8486993	Aug 30, 2024	DP U-1752			
	8586084	Aug 30, 2024	U-1752			
	8591946	Aug 30, 2024	DP			
	8617599	Aug 30, 2024	DP			
	8623410	Aug 30, 2024	DP			
	8623411	Aug 30, 2024	U-1752			
	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			
	8889185	Aug 30, 2024	U-1752			
	8889186	Aug 30, 2024	U-1752			
	9161907	Aug 30, 2024	DP U-1752			
	9549918	May 30, 2028	DP			
	9757362	Aug 30, 2024	DP			
	9763920	Aug 30, 2024	DP			
<u>TACROLIMUS - PROGRAF</u>						
N 210115 001					ODE-269	May 24, 2025
<u>TACROLIMUS - PROGRAF</u>						
N 210115 002					ODE-269	May 24, 2025
<u>TADALAFIL - CIALIS</u>						
N 021368 001	6943166	Apr 26, 2020	U-1184		M-219	Feb 15, 2021
	6943166	Apr 26, 2020	U-155		PED	Aug 15, 2021
	6943166	Apr 26, 2020	U-614			
	6943166*PED	Oct 26, 2020				
<u>TADALAFIL - CIALIS</u>						
N 021368 002	6943166	Apr 26, 2020	U-155		M-219	Feb 15, 2021
	6943166	Apr 26, 2020	U-614		PED	Aug 15, 2021
	6943166*PED	Oct 26, 2020				

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APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>TADALAFIL - CIALIS</u>						
N 021368 002	6943166	Apr 26, 2020	U-155		M-219	Feb 15, 2021
	6943166	Apr 26, 2020	U-614		PED	Aug 15, 2021
	6943166*PED	Oct 26, 2020				
<u>TADALAFIL - CIALIS</u>						
N 021368 003	6943166	Apr 26, 2020	U-614		M-219	Feb 15, 2021
	6943166*PED	Oct 26, 2020			PED	Aug 15, 2021
<u>TADALAFIL - CIALIS</u>						
N 021368 004	6943166	Apr 26, 2020	U-155		M-219	Feb 15, 2021
	6943166*PED	Oct 26, 2020			PED	Aug 15, 2021
<u>TAFAMIDIS - VYNDAMAX</u>						
N 212161 001	7214695	Apr 27, 2024	DS DP		NCE	May 03, 2024
	7214696	Dec 19, 2023		U-2524	ODE-237	May 03, 2026
	9770441	Aug 31, 2035	DS DP	U-2524		
<u>TAFAMIDIS MEGLUMINE - VYNDAQEL</u>						
N 211996 001	7214695	Apr 27, 2024	DS DP		NCE	May 03, 2024
	7214696	Dec 19, 2023		U-2524	ODE-237	May 03, 2026
	8168663	Dec 19, 2023	DS DP			
	8653119	Jan 28, 2024		U-2524		
<u>TAFENOQUINE SUCCINATE - ARAKODA</u>						
N 210607 001	10342791	Dec 02, 2035		U-2582	NCE NP	Jul 20, 2023 Aug 08, 2021
<u>TAFENOQUINE SUCCINATE - KRINTAFEL</u>						
N 210795 001					NCE ODE-201	Jul 20, 2023 Jul 20, 2025
<u>TAFLUPROST - ZIOPATAN</u>						
N 202514 001	5886035	Dec 18, 2022	DS DP	U-778		
	9999593	May 28, 2029		DP		
<u>TALAZOPARIB TOSYLATE - TALZENNA</u>						
N 211651 001	10189837	Oct 20, 2031	DS DP		NCE	Oct 16, 2023
	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 002	10189837	Oct 20, 2031	DS DP		NCE	Oct 16, 2023
	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>TALC - STERITALC</u>						
N 205555 001					ODE-143 ODE-191	May 01, 2024 May 01, 2024
<u>TALC - STERITALC</u>						
N 205555 002					ODE-143 ODE-191	May 01, 2024 May 01, 2024
<u>TALC - STERITALC</u>						
N 205555 003					ODE-143 ODE-191	May 01, 2024 May 01, 2024
<u>TALIGLUCERASE ALFA - ELELYSO</u>						
N 022458 001	8227230	Feb 24, 2024	DS DP			
	8741620	Feb 24, 2024	DS DP			
	8790641	Oct 18, 2025		U-1564		
	8790641	Oct 18, 2025		U-1574		
<u>TAPENTADOL HYDROCHLORIDE - NUCYNTA</u>						
N 022304 001	7994364	Jun 27, 2025	DS DP	U-931		
	RE39593	Aug 05, 2022	DS DP	U-931		

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<u>TAPENTADOL HYDROCHLORIDE - NUCYNTA</u>						
N 022304	002	7994364	Jun 27, 2025	DS DP U-931		
		RE39593	Aug 05, 2022	DS DP U-931		
<u>TAPENTADOL HYDROCHLORIDE - NUCYNTA</u>						
N 022304	003	7994364	Jun 27, 2025	DS DP U-931		
		RE39593	Aug 05, 2022	DS DP U-931		
<u>TAPENTADOL HYDROCHLORIDE - NUCYNTA ER</u>						
N 200533	001	7994364	Jun 27, 2025	DS DP U-1178		
		7994364	Jun 27, 2025	DS DP U-1276		
		8075872	Nov 20, 2023	DP		
		8114383	Oct 10, 2024	DP	Y	
		8309060	Nov 20, 2023	DP U-1178		
		8309060	Nov 20, 2023	DP U-1276		
		8420056	Nov 20, 2023	DP		
		8536130	Sep 22, 2028	U-1276		
		RE39593	Aug 05, 2022	DS DP U-1178		
		RE39593	Aug 05, 2022	DS DP U-1276		
<u>TAPENTADOL HYDROCHLORIDE - NUCYNTA ER</u>						
N 200533	002	7994364	Jun 27, 2025	DS DP U-1178		
		7994364	Jun 27, 2025	DS DP U-1276		
		8075872	Nov 20, 2023	DP		
		8114383	Oct 10, 2024	DP	Y	
		8309060	Nov 20, 2023	DP U-1178		
		8309060	Nov 20, 2023	DP U-1276		
		8420056	Nov 20, 2023	DP		
		8536130	Sep 22, 2028	U-1276		
		RE39593	Aug 05, 2022	DS DP U-1178		
		RE39593	Aug 05, 2022	DS DP U-1276		
<u>TAPENTADOL HYDROCHLORIDE - NUCYNTA ER</u>						
N 200533	003	7994364	Jun 27, 2025	DS DP U-1178		
		7994364	Jun 27, 2025	DS DP U-1276		
		8075872	Nov 20, 2023	DP		
		8114383	Oct 10, 2024	DP	Y	
		8309060	Nov 20, 2023	DP U-1178		
		8309060	Nov 20, 2023	DP U-1276		
		8420056	Nov 20, 2023	DP		
		8536130	Sep 22, 2028	U-1276		
		RE39593	Aug 05, 2022	DS DP U-1178		
		RE39593	Aug 05, 2022	DS DP U-1276		
<u>TAPENTADOL HYDROCHLORIDE - NUCYNTA ER</u>						
N 200533	004	7994364	Jun 27, 2025	DS DP U-1178		
		7994364	Jun 27, 2025	DS DP U-1276		
		8075872	Nov 20, 2023	DP		
		8114383	Oct 10, 2024	DP	Y	
		8309060	Nov 20, 2023	DP U-1178		
		8309060	Nov 20, 2023	DP U-1276		
		8420056	Nov 20, 2023	DP		
		8536130	Sep 22, 2028	U-1276		
		RE39593	Aug 05, 2022	DS DP U-1178		
		RE39593	Aug 05, 2022	DS DP U-1276		
<u>TAPENTADOL HYDROCHLORIDE - NUCYNTA ER</u>						
N 200533	005	7994364	Jun 27, 2025	DS DP U-1178		
		7994364	Jun 27, 2025	DS DP U-1276		
		8075872	Nov 20, 2023	DP		
		8114383	Oct 10, 2024	DP	Y	
		8309060	Nov 20, 2023	DP U-1178		
		8309060	Nov 20, 2023	DP U-1276		
		8420056	Nov 20, 2023	DP		
		8536130	Sep 22, 2028	U-1276		
		RE39593	Aug 05, 2022	DS DP U-1178		
		RE39593	Aug 05, 2022	DS DP U-1276		
<u>TAPENTADOL HYDROCHLORIDE - NUCYNTA</u>						
N 203794	001	7994364	Jun 27, 2025	DS DP U-1289		
		RE39593	Aug 05, 2022	DS DP U-1289		

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<u>TAPENTADOL HYDROCHLORIDE - NUCYNTA</u>						
N 203794	001 7994364	Jun 27, 2025	DS DP U-1289			
	RE39593	Aug 05, 2022	DS DP U-1289			
<u>TASIMELTEON - HETLIOZ</u>						
N 205677	001 10071977	Feb 12, 2035	DS DP		ODE-59	Jan 31, 2021
	10149829	Jan 25, 2033	U-2477			
	10376487	Jul 27, 2035	U-2615			
	10449176	Jan 25, 2033	U-2149			
	5856529	Dec 09, 2022	DS DP U-2149			
	9060995	Jan 25, 2033	U-1710			
	9539234	Jan 25, 2033	U-1934			
	9549913	Jan 25, 2033	U-1486			
	9730910	May 17, 2034	U-2085			
	9855241	Jan 25, 2033	U-2149			
	RE46604	Jan 25, 2033	U-2147			
<u>TAVABOROLE - KERYDIN</u>						
N 204427	001 7582621	May 26, 2027	U-2016	Y		
	7582621*PED	Nov 26, 2027				
	9549938	Feb 16, 2026	U-1951			
	9549938*PED	Aug 16, 2026				
	9566289	Feb 16, 2026	DP			
	9566289*PED	Aug 16, 2026				
	9566290	Feb 16, 2026	U-1970			
	9566290*PED	Aug 16, 2026				
	9572823	Feb 16, 2026	U-1970			
	9572823*PED	Aug 16, 2026				
<u>TAZAROTENE - FABIOR</u>						
N 202428	001 8808716	Feb 24, 2030	DP			
<u>TAZAROTENE - ARAZLO</u>						
N 211882	001 6517847	Aug 03, 2020	DP U-2368		NP	Dec 18, 2022
<u>TAZEMETOSTAT HYDROBROMIDE - TAZVERIK</u>						
N 211723	001 10155002	Apr 13, 2032	U-2736		NCE	Jan 23, 2025
	10245269	Apr 11, 2033	U-2737			
	10369155	Oct 16, 2035	U-2736			
	10420775	Apr 13, 2032	U-2736			
	8410088	Apr 13, 2032	DS DP			
	9090562	Apr 13, 2032	DS DP			
	9394283	Apr 11, 2033	DS DP			
	9522152	Apr 13, 2032	U-2738			
	9549931	Apr 13, 2032	U-2736			
	9688665	Aug 22, 2034	U-2736			
	9855275	Apr 13, 2032	U-2736			
	9872862	Apr 11, 2033	U-2738			
	9889138	Oct 16, 2035	U-2736			
<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, 2020	DS		ODE-67	Jun 13, 2021
	9439985	Sep 27, 2033	DS DP			
<u>TECOVIRIMAT - TPOXX</u>						
N 208627	001 7737168	May 03, 2027	U-2346		NCE	Jul 13, 2023
	8039504	Jul 23, 2027	DP		ODE-200	Jul 13, 2025
	8124643	Jun 18, 2024	DS DP			
	8530509	Jun 18, 2024	DP			
	8802714	Jun 18, 2024	U-2346			
	9339466	Mar 23, 2031	DS DP			
<u>TEDIZOLID PHOSPHATE - SIVEXTRO</u>						
N 205435	001 10065947	Feb 03, 2030	DP		NCE	Jun 20, 2019
	10442829	Feb 03, 2030	DS		GAIN	Jun 20, 2024
	7816379	Apr 20, 2026	DS DP U-2507			
	7816379	Apr 20, 2026	DS DP U-282			
	8420676	Feb 23, 2028	DS DP U-282			

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<u>TEDIZOLID PHOSPHATE - SIVEXTRO</u>						
N 205435	001 8426389	Dec 31, 2030	DS DP U-282			
	9624250	Feb 03, 2030	DS DP U-2507			
	9988406	Feb 03, 2030	DP			
<u>TEDIZOLID PHOSPHATE - SIVEXTRO</u>						
N 205436	001 10065947	Feb 03, 2030	DP		NCE	Jun 20, 2019
	10442829	Feb 03, 2030	DS		GAIN	Jun 20, 2024
	7816379	Apr 20, 2026	DS DP U-2507			
	7816379	Apr 20, 2026	DS DP U-282			
	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 RECOMBINANT - GATTEX KIT</u>						
N 203441	001 5789379	Apr 14, 2020	DS DP U-1320		ODE-240	May 16, 2026
	5789379*PED	Oct 14, 2020				
	7056886	Sep 18, 2022	DP U-1320			
	7056886*PED	Mar 18, 2023				
	7847061	Nov 01, 2025	U-1320			
	7847061*PED	May 01, 2026				
	9060992	Nov 01, 2025	U-1320			
	9060992*PED	May 01, 2026				
	9539310	Nov 01, 2025	U-1320			
	9539310*PED	May 01, 2026				
	9545434	Nov 01, 2025	U-1320			
	9545434*PED	May 01, 2026				
	9545435	Nov 01, 2025	U-1320			
	9545435*PED	May 01, 2026				
	9555079	Nov 01, 2025	U-1320			
	9555079*PED	May 01, 2026				
	9572867	Nov 01, 2025	U-1320			
	9572867*PED	May 01, 2026				
	9592273	Nov 01, 2025	U-1320			
	9592273*PED	May 01, 2026				
	9592274	Nov 01, 2025	U-1320			
	9592274*PED	May 01, 2026				
	9968655	Nov 01, 2025	U-2308			
	9968655*PED	May 01, 2026				
	9968656	Nov 01, 2025	U-2308			
	9968656*PED	May 01, 2026				
	9968658	Nov 01, 2025	U-1320			
	9968658*PED	May 01, 2026				
	9974835	Nov 01, 2025	U-1320			
	9974835*PED	May 01, 2026				
	9974837	Nov 01, 2025	U-1320			
	9974837*PED	May 01, 2026				
	9981014	Nov 01, 2025	U-1320			
	9981014*PED	May 01, 2026				
	9981016	Nov 01, 2025	U-1320			
	9981016*PED	May 01, 2026				
	9987334	Nov 01, 2025	U-1320			
	9987334*PED	May 01, 2026				
	9987335	Nov 01, 2025	U-1320			
	9987335*PED	May 01, 2026				
	9993528	Nov 01, 2025	U-1320			
	9993528*PED	May 01, 2026				
<u>TELAPREVIR - INCIVEK</u>						
N 201917	001 7820671	Feb 25, 2025	DS DP			
	8431615	May 30, 2028	U-1398			
	8529882	Aug 31, 2021	U-1398			
<u>TELAVANCIN HYDROCHLORIDE - VIBATIV</u>						
N 022110	001 6635618	Sep 11, 2023	DS DP U-728			
	6858584	Aug 24, 2022	DP			
	6872701	Jun 05, 2021	DP			
	7008923	May 06, 2021	U-1005			
	7208471	May 01, 2021	DS DP			
	7351691	May 01, 2021	DS DP U-728			

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<u>TELAVANCIN HYDROCHLORIDE - VIBATIV</u>						
N 022110 001	7531623	Jan 01, 2027	DS			
	7544364	May 01, 2021	DP			
	7700550	May 01, 2021		U-282		
	8101575	May 01, 2021	DP			
	8158580	May 01, 2021	DP			
<u>TELAVANCIN HYDROCHLORIDE - VIBATIV</u>						
N 022110 002	6635618	Sep 11, 2023	DS DP	U-728		
	6858584	Aug 24, 2022	DP			
	6872701	Jun 05, 2021	DP			
	7008923	May 06, 2021		U-1005		
	7208471	May 01, 2021	DS DP			
	7351691	May 01, 2021	DS DP	U-728		
	7531623	Jan 01, 2027	DS			
	7544364	May 01, 2021	DP			
	7700550	May 01, 2021		U-282		
	8101575	May 01, 2021	DP			
	8158580	May 01, 2021	DP			
<u>TELBIVUDINE - TYZEKA</u>						
N 022011 001	6569837	Oct 25, 2020		U-782		
	6569837	Oct 25, 2020		U-999		
	7589079	Sep 11, 2023	DS DP	U-999		
	7858594	Sep 11, 2023	DS DP	U-999		
<u>TELBIVUDINE - TYZEKA</u>						
N 022154 001	6569837	Oct 25, 2020		U-999		
	7858594	Sep 11, 2023	DS DP	U-999		
<u>TELMISARTAN - MICARDIS</u>						
N 020850 001	6358986	Jan 10, 2020				
<u>TELMISARTAN - MICARDIS</u>						
N 020850 002	6358986	Jan 10, 2020				
	7998953	Jun 06, 2020		U-1177		
	8003679	Oct 06, 2022		U-1176		
<u>TELMISARTAN - MICARDIS</u>						
N 020850 003	6358986	Jan 10, 2020				
<u>TELOTTRISTAT ETIPRATE - XERMELO</u>						
N 208794 001	7553840	Dec 11, 2027	DS		NCE	Feb 28, 2022
	7709493	Dec 11, 2027	DS	U-1979	ODE-132	Feb 28, 2024
	7968559	Dec 11, 2027		U-1979		
	8193204	Feb 27, 2031	DS			
	8653094	Dec 19, 2028		U-1979		
<u>TEMOZOLOMIDE - TEMODAR</u>						
N 022277 001	6987108	Sep 08, 2023	DP			
	7786118	Feb 21, 2023	DP			
	8623868	Feb 21, 2023	DP			
<u>TEMSIROLIMUS - TORISEL</u>						
N 022088 001	8026276	Jan 20, 2026	DP			
	8299116	Jul 25, 2023	DP			
	8455539	Jul 25, 2023	DP			
	8455539*PED	Jan 25, 2024				
	8722700	Jul 25, 2023	DP			
	8722700*PED	Jan 25, 2024				
	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	8541448	Dec 30, 2029	DS DP		NCE	Sep 12, 2024
	8969377	Dec 30, 2029	DS DP			
	9006281	May 02, 2030		U-2626		
	9408840	Dec 30, 2029		U-2626		

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<u>TENOFOVIR ALAFENAMIDE FUMARATE - VEMLIDY</u>						
N 208464 001	7390791	May 07, 2022	DS DP		M-255	Feb 04, 2023
	7803788	Feb 02, 2022		U-999	NCE	Nov 05, 2020
	8754065	Aug 15, 2032	DS DP	U-999		
	9296769	Aug 15, 2032	DS DP	U-999		
<u>TENOFOVIR DISOPROXIL FUMARATE - VIREAD</u>						
N 021356 001					NPP PED	Dec 11, 2021 Jun 11, 2022
<u>TENOFOVIR DISOPROXIL FUMARATE - VIREAD</u>						
N 021356 002					NPP PED	Dec 11, 2021 Jun 11, 2022
<u>TENOFOVIR DISOPROXIL FUMARATE - VIREAD</u>						
N 021356 003					NPP PED	Dec 11, 2021 Jun 11, 2022
<u>TENOFOVIR DISOPROXIL FUMARATE - VIREAD</u>						
N 021356 004					NPP PED	Dec 11, 2021 Jun 11, 2022
<u>TENOFOVIR DISOPROXIL FUMARATE - VIREAD</u>						
N 022577 001					NPP PED	Dec 11, 2021 Jun 11, 2022
<u>TERIFLUNOMIDE - AUBAGIO</u>						
N 202992 001	6794410	Sep 12, 2026		U-1285		
	8802735	Sep 14, 2030	DP			
	9186346	Feb 04, 2034		U-1786		
<u>TERIFLUNOMIDE - AUBAGIO</u>						
N 202992 002	6794410	Sep 12, 2026		U-1285		
	8802735	Sep 14, 2030	DP			
	9186346	Feb 04, 2034		U-1786		
<u>TERIPARATIDE - FORTEO</u>						
N 021318 001	7517334	Mar 25, 2025		DP		
<u>TERIPARATIDE - FORTEO</u>						
N 021318 002	7517334	Mar 25, 2025		DP		
<u>TESAMORELIN ACETATE - EGRIFTA</u>						
N 022505 001	5861379	May 26, 2020	DS DP	U-1100		
	7144577	Jul 14, 2020		U-1100		
	7316997	Aug 14, 2023		U-1100		
	8314066	Aug 14, 2023		U-1100		
	8435945	Aug 14, 2023		U-1100		
<u>TESAMORELIN ACETATE - EGRIFTA</u>						
N 022505 002	5861379	May 26, 2020	DS DP	U-1100		
	7144577	Jul 14, 2020		U-1100		
	7316997	Aug 14, 2023		U-1100		
	8314066	Aug 14, 2023		U-1100		
	8435945	Aug 14, 2023		U-1100		
<u>TESTOSTERONE - ANDROGEL</u>						
N 021015 001	6503894	Aug 30, 2020		U-490		
	9125816	Aug 30, 2020		U-490		
	9125816*PED	Mar 02, 2021				
	9132089	Aug 30, 2020		U-490		
	9132089*PED	Mar 02, 2021				
<u>TESTOSTERONE - ANDROGEL</u>						
N 021015 002	6503894	Aug 30, 2020		U-490		
	9125816	Aug 30, 2020		U-490		
	9125816*PED	Mar 02, 2021				
	9132089	Aug 30, 2020		U-490		
	9132089*PED	Mar 02, 2021				

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<u>TESTOSTERONE - ANDROGEL</u>						
N 021015	003	6503894	Aug 30, 2020	U-490		
		9125816	Aug 30, 2020	U-490		
		9125816*PED	Mar 02, 2021			
		9132089	Aug 30, 2020	U-490		
		9132089*PED	Mar 02, 2021			
<u>TESTOSTERONE - TESTIM</u>						
N 021454	001	7320968	Jan 18, 2025	U-843		
		7608605	Apr 21, 2023	U-1009		
		7608606	Apr 21, 2023	U-1009		
		7608607	Apr 21, 2023	U-1009		
		7608608	Apr 21, 2023	U-1009		
		7608609	Apr 21, 2023	U-1009		
		7608610	Apr 21, 2023	U-1009		
		7935690	Apr 21, 2023	U-1009		
		8063029	Apr 21, 2023	U-843		
		8178518	Apr 21, 2023	DP		
<u>TESTOSTERONE - ANDROGEL</u>						
N 022309	001	6503894	Aug 30, 2020	U-1103		
		6503894*PED	Mar 02, 2021			
		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		
		9125816	Aug 30, 2020	U-1103		
		9125816*PED	Mar 02, 2021			
		9132089	Aug 30, 2020	U-1103		
		9132089*PED	Mar 02, 2021			
<u>TESTOSTERONE - ANDROGEL</u>						
N 022309	002	6503894	Aug 30, 2020	U-1103		
		6503894*PED	Mar 02, 2021			
		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		
		9125816	Aug 30, 2020	U-1103		
		9125816*PED	Mar 02, 2021			
		9132089	Aug 30, 2020	U-1103		
		9132089*PED	Mar 02, 2021			
<u>TESTOSTERONE - ANDROGEL</u>						
N 022309	003	6503894	Aug 30, 2020	U-1103		
		6503894*PED	Mar 02, 2021			
		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		
		9125816	Aug 30, 2020	U-1103		
		9125816*PED	Mar 02, 2021			
		9132089	Aug 30, 2020	U-1103		
		9132089*PED	Mar 02, 2021			
<u>TESTOSTERONE - AXIRON</u>						
N 022504	001	8419307	Feb 26, 2027	U-1386		
		8435944	Sep 27, 2027	U-1390		
		8784878	Jul 13, 2023	DP U-1545		

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<u>TESTOSTERONE - AXIRON</u>						
N 022504	001	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		
<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	8574622	Feb 04, 2024	DP		
		8784869	Feb 04, 2024	DP		
		8784882	Feb 04, 2024	DP U-1557		
		8877230	Feb 04, 2024	U-1616		
<u>TESTOSTERONE ENANTHATE - XYOSTED (AUTOINJECTOR)</u>						
N 209863	001	8021335	Oct 04, 2026	DP	NP	Sep 28, 2021
		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	8021335	Oct 04, 2026	DP	NP	Sep 28, 2021
		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	8021335	Oct 04, 2026	DP	NP	Sep 28, 2021
		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	Feb 27, 2026	U-1500		
<u>TESTOSTERONE UNDECANOATE - JATENZO</u>						
N 206089	001	10543219	Apr 12, 2030	U-2506	NP	Mar 27, 2022
		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	NP	Mar 27, 2022
		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	NP	Mar 27, 2022
		8241664	Mar 29, 2029	DP U-2506		
		8492369	Dec 20, 2030	DP U-2506		
		8778916	Apr 12, 2030	DP		

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<u>THALIDOMIDE - THALOMID</u>						
N 020785 001	6315720	Oct 23, 2020	U-442			
	6315720	Oct 23, 2020	U-731			
	6561977	Oct 23, 2020	U-371			
	6561977	Oct 23, 2020	U-731			
	6755784	Oct 23, 2020	U-371			
	6755784	Oct 23, 2020	U-731			
	6869399	Oct 23, 2020	U-371			
	6869399	Oct 23, 2020	U-731			
	6869399	Oct 23, 2020	U-732			
	6869399	Oct 23, 2020	U-733			
	7141018	Oct 23, 2020	U-371			
	7141018	Oct 23, 2020	U-731			
	7141018	Oct 23, 2020	U-732			
	7141018	Oct 23, 2020	U-733			
	7230012	Dec 09, 2023		DP		
	7959566	Oct 23, 2020	U-1155			
	8315886	Oct 23, 2020	U-1249			
	8626531	Oct 23, 2020	U-1465			
<u>THALIDOMIDE - THALOMID</u>						
N 020785 002	6315720	Oct 23, 2020	U-442			
	6315720	Oct 23, 2020	U-731			
	6561977	Oct 23, 2020	U-371			
	6561977	Oct 23, 2020	U-731			
	6755784	Oct 23, 2020	U-371			
	6755784	Oct 23, 2020	U-731			
	6869399	Oct 23, 2020	U-371			
	6869399	Oct 23, 2020	U-731			
	6869399	Oct 23, 2020	U-732			
	6869399	Oct 23, 2020	U-733			
	7141018	Oct 23, 2020	U-371			
	7141018	Oct 23, 2020	U-731			
	7141018	Oct 23, 2020	U-732			
	7141018	Oct 23, 2020	U-733			
	7230012	Dec 09, 2023		DP		
	7959566	Oct 23, 2020	U-1155			
	8315886	Oct 23, 2020	U-1249			
	8626531	Oct 23, 2020	U-1465			
<u>THALIDOMIDE - THALOMID</u>						
N 020785 003	6315720	Oct 23, 2020	U-442			
	6315720	Oct 23, 2020	U-731			
	6561977	Oct 23, 2020	U-371			
	6561977	Oct 23, 2020	U-731			
	6755784	Oct 23, 2020	U-371			
	6755784	Oct 23, 2020	U-731			
	6869399	Oct 23, 2020	U-371			
	6869399	Oct 23, 2020	U-731			
	6869399	Oct 23, 2020	U-732			
	6869399	Oct 23, 2020	U-733			
	7141018	Oct 23, 2020	U-371			
	7141018	Oct 23, 2020	U-731			
	7141018	Oct 23, 2020	U-732			
	7141018	Oct 23, 2020	U-733			
	7230012	Dec 09, 2023		DP		
	7959566	Oct 23, 2020	U-1155			
	8315886	Oct 23, 2020	U-1249			
	8626531	Oct 23, 2020	U-1465			
<u>THALIDOMIDE - THALOMID</u>						
N 020785 004	6315720	Oct 23, 2020	U-731			
	6561977	Oct 23, 2020	U-731			
	6755784	Oct 23, 2020	U-731			
	6869399	Oct 23, 2020	U-731			
	7141018	Oct 23, 2020	U-731			
	7959566	Oct 23, 2020	U-1155			
	8315886	Oct 23, 2020	U-1249			
	8626531	Oct 23, 2020	U-1465			

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<u>THIOTEPA - TEPADINA</u>						
N 208264	001				I-747 ODE-129	Jan 26, 2020 Jan 26, 2024
<u>THIOTEPA - TEPADINA</u>						
N 208264	002				I-747 ODE-129	Jan 26, 2020 Jan 26, 2024
<u>TICAGRELOR - BRILINTA</u>						
N 022433	001	10300065	Jan 27, 2036	U-2541		
		10300065	Jan 27, 2036	U-2542		
		7265124	Jul 09, 2021	DS DP U-1171		
		7265124	Jul 09, 2021	DS DP U-1860		
		7265124	Jul 09, 2021	DS DP U-1868		
		7265124	Jul 09, 2021	DS DP U-1869		
		8425934	Apr 17, 2030	DP		
		RE46276	Oct 30, 2024	DS DP U-1935		
		RE46276	Oct 30, 2024	DS DP U-1936		
		RE46276	Oct 30, 2024	DS DP U-1937		
		RE46276	Oct 30, 2024	DS DP U-1938		
<u>TICAGRELOR - BRILINTA</u>						
N 022433	002	10300065	Jan 27, 2036	U-2541		
		10300065	Jan 27, 2036	U-2542		
		7265124	Jul 09, 2021	DS DP U-1171		
		7265124	Jul 09, 2021	DS DP U-1860		
		7265124	Jul 09, 2021	DS DP U-1868		
		7265124	Jul 09, 2021	DS DP U-1869		
		8425934	Apr 17, 2030	DP		
		RE46276	Oct 30, 2024	DS DP U-1935		
		RE46276	Oct 30, 2024	DS DP U-1936		
		RE46276	Oct 30, 2024	DS DP U-1937		
		RE46276	Oct 30, 2024	DS DP U-1938		
<u>TIGECYCLINE - TYGACIL</u>						
N 021821	001	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>TIOTROPIUM BROMIDE - SPIRIVA</u>						
N 021395	001	6777423	Sep 24, 2021	DS DP		
		6777423*PED	Mar 24, 2022			
		6908928	Sep 24, 2021	DS DP U-566		
		6908928	Sep 24, 2021	DS DP U-762		
		6908928*PED	Mar 24, 2022			
		7070800	Jan 22, 2022	DP U-566		
		7070800*PED	Jul 22, 2022			
		7309707	Sep 24, 2021	DS DP		
		7309707*PED	Mar 24, 2022			
		7642268	Sep 24, 2021	DS DP		
		7642268*PED	Mar 24, 2022			
		7694676	Mar 12, 2027	DP		
		7694676*PED	Sep 12, 2027			
		8022082	Jan 19, 2026	DP U-1186		
		8022082*PED	Jul 19, 2026			
		9010323	Apr 19, 2030	DP		
		RE38912	Oct 11, 2021	DP		
		RE38912*PED	Apr 11, 2022			
<u>TIOTROPIUM BROMIDE - SPIRIVA RESPIMAT</u>						
N 021936	001	6988496	Feb 23, 2020	DP	NPP	Feb 15, 2020
		6988496*PED	Aug 23, 2020		PED	Aug 15, 2020
		7284474	Aug 26, 2024	DP		

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<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				
	7988001	Aug 04, 2021	DP			
	7988001*PED	Feb 04, 2022				
	8733341	Oct 16, 2030	DP			
	8733341*PED	Apr 16, 2031				
	9027967	Mar 31, 2027	DP			
	9027967*PED	Oct 01, 2027				
<u>TIOTROPIUM BROMIDE - SPIRIVA RESPIMAT</u>						
N 021936 002	6988496	Feb 23, 2020	DP		NPP	Feb 15, 2020
	6988496*PED	Aug 23, 2020			PED	Aug 15, 2020
	7284474	Aug 26, 2024	DP			
	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				
	7988001	Aug 04, 2021	DP			
	7988001*PED	Feb 04, 2022				
	8733341	Oct 16, 2030	DP			
	8733341*PED	Apr 16, 2031				
	9027967	Mar 31, 2027	DP			
	9027967*PED	Oct 01, 2027				
<u>TIPIRACIL HYDROCHLORIDE; TRIFLURIDINE - LONSURE</u>						
N 207981 001	10456399	Feb 03, 2037		U-2642	I-794	Feb 22, 2022
	10457666	Jun 17, 2034	DS DP		NCE	Sep 22, 2020
	6479500	Mar 16, 2020		U-1751	ODE-229	Feb 22, 2026
	9527833	Jun 17, 2034	DS DP			
	RE46284	Dec 16, 2026		U-1751		
	RE46284	Dec 16, 2026		U-2503		
<u>TIPIRACIL HYDROCHLORIDE; TRIFLURIDINE - LONSURE</u>						
N 207981 002	10456399	Feb 03, 2037		U-2642	I-794	Feb 22, 2022
	10457666	Jun 17, 2034	DS DP		NCE	Sep 22, 2020
	6479500	Mar 16, 2020		U-1751	ODE-229	Feb 22, 2026
	9527833	Jun 17, 2034	DS DP			
	RE46284	Dec 16, 2026		U-1751		
	RE46284	Dec 16, 2026		U-2503		
<u>TIROFIBAN HYDROCHLORIDE - AGGRASTAT</u>						
N 020912 001	6770660	May 01, 2023		U-1444		
<u>TIROFIBAN HYDROCHLORIDE - AGGRASTAT</u>						
N 020912 002	6770660	May 01, 2023		U-1444		
<u>TIROFIBAN HYDROCHLORIDE - AGGRASTAT</u>						
N 020913 001	6770660	May 01, 2023		U-1444		
<u>TIROFIBAN HYDROCHLORIDE - AGGRASTAT</u>						
N 020913 002	6770660	May 01, 2023		U-1444		
<u>TIROFIBAN HYDROCHLORIDE - AGGRASTAT</u>						
N 020913 003	6770660	May 01, 2023		U-1444		
<u>TOBRAMYCIN - TOBI PODHALER</u>						
N 201688 001	10207066	Nov 04, 2030	DP	U-909		
	7368102	Dec 19, 2022	DP	U-909		
	7442388	May 10, 2020	DP			
	7516741	Jan 11, 2024	DP			
	7559325	Oct 27, 2025	DP			
	8069851	Sep 24, 2024	DP			

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<u>TOBRAMYCIN - TOBI PODHALER</u>						
N 201688	001	8349294	May 10, 2020	DP		
		8664187	Jun 20, 2025	U-909		
		8715623	Dec 19, 2022	DP U-909		
		8869794	Sep 12, 2028	DP U-909		
		9421166	Dec 19, 2022	DP U-909		
<u>TOBRAMYCIN - BETHKIS</u>						
N 201820	001	6987094	Sep 22, 2022	DP		
		7696178	Sep 22, 2022	DP		
		7939502	Jun 14, 2022	U-1324		
<u>TOFACITINIB CITRATE - XELJANZ</u>						
N 203214	001	6956041	Dec 08, 2020	DP	I-761	Dec 14, 2020
		6965027	Mar 25, 2023	DS	I-780	May 30, 2021
		7091208	Dec 08, 2020	U-247		
		7265221	Dec 08, 2020	DS		
		7301023	May 23, 2022	DS		
		7842699	Dec 08, 2020	U-2322		
		RE41783	Dec 08, 2025	DS		
<u>TOFACITINIB CITRATE - XELJANZ</u>						
N 203214	002	6956041	Dec 08, 2020	DP	I-780	May 30, 2021
		6965027	Mar 25, 2023	DS		
		7265221	Dec 08, 2020	DS		
		7301023	May 23, 2022	DS		
		7842699	Dec 08, 2020	U-2322		
		RE41783	Dec 08, 2025	DS		
<u>TOFACITINIB CITRATE - XELJANZ XR</u>						
N 208246	001	6956041	Dec 08, 2020	DP	I-761	Dec 14, 2020
		6965027	Mar 25, 2023	DS		
		7091208	Dec 08, 2020	U-247		
		7265221	Dec 08, 2020	DS		
		7301023	May 23, 2022	DS		
		7842699	Dec 08, 2020	U-2322		
		9937181	Mar 14, 2034	DP		
		RE41783	Dec 08, 2025	DS		
<u>TOFACITINIB CITRATE - XELJANZ XR</u>						
N 208246	002	6956041	Dec 08, 2020	DP		
		6965027	Mar 25, 2023	DS		
		7265221	Dec 08, 2020	DS		
		7301023	May 23, 2022	DS		
		7842699	Dec 08, 2020	U-2322		
		RE41783	Dec 08, 2025	DS		
<u>TOLTERODINE TARTRATE - DETROL LA</u>						
N 021228	001	6911217*PED	May 11, 2020			
<u>TOLTERODINE TARTRATE - DETROL LA</u>						
N 021228	002	6911217*PED	May 11, 2020			
<u>TOLVAPTAN - SAMSCA</u>						
N 022275	001	5753677	May 19, 2020	U-978		
		8501730	Sep 01, 2026	DS		
<u>TOLVAPTAN - SAMSCA</u>						
N 022275	002	5753677	May 19, 2020	U-978		
		8501730	Sep 01, 2026	DS		
<u>TOLVAPTAN - SAMSCA</u>						
N 022275	003	5753677	May 19, 2020	U-978		
		8501730	Sep 01, 2026	DS		
<u>TOLVAPTAN - JYNARQUE</u>						
N 204441	001	5753677	May 19, 2020	U-2307	I-779	Apr 23, 2021
		8501730	Sep 01, 2026	DS	ODE-178	Apr 23, 2025

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<u>TOLVAPTAN - JYNARQUE</u>						
N 204441 002	5753677	May 19, 2020	U-2307		I-779	Apr 23, 2021
	8501730	Sep 01, 2026	DS		ODE-178	Apr 23, 2025
<u>TOLVAPTAN - JYNARQUE</u>						
N 204441 003	5753677	May 19, 2020	U-2307		I-779	Apr 23, 2021
	8501730	Sep 01, 2026	DS		ODE-178	Apr 23, 2025
<u>TOLVAPTAN - JYNARQUE</u>						
N 204441 004	5753677	May 19, 2020	U-2307		I-779	Apr 23, 2021
	8501730	Sep 01, 2026	DS		ODE-178	Apr 23, 2025
<u>TOLVAPTAN - JYNARQUE</u>						
N 204441 005	5753677	May 19, 2020	U-2307		I-779	Apr 23, 2021
	8501730	Sep 01, 2026	DS		ODE-178	Apr 23, 2025
<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			
	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			

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<u>TOPIRAMATE - TROKENDI XR</u>						
N 201635	003	8889191				Nov 16, 2027
		8889191				Nov 16, 2027
		8992989		DP		Nov 16, 2027
		8992989		DP		Nov 16, 2027
		9549940		DP		Nov 16, 2027
		9549940		DP		Nov 16, 2027
		9555004		DP		Nov 16, 2027
		9555004		DP		Nov 16, 2027
		9622983		DP		Nov 16, 2027
		9622983		DP		Nov 16, 2027
<u>TOPIRAMATE - TROKENDI XR</u>						
N 201635	004	10314790		DP		Nov 16, 2027
		10314790		DP		Nov 16, 2027
		8298576		DP		Apr 04, 2028
		8298576		DP		Apr 04, 2028
		8298580		DP		Nov 16, 2027
		8298580		DP		Nov 16, 2027
		8663683		DP		Nov 16, 2027
		8663683		DP		Nov 16, 2027
		8877248		DP		Nov 16, 2027
		8877248		DP		Nov 16, 2027
		8889191				Nov 16, 2027
		8889191				Nov 16, 2027
		8992989		DP		Nov 16, 2027
		8992989		DP		Nov 16, 2027
		9549940		DP		Nov 16, 2027
		9549940		DP		Nov 16, 2027
		9555004		DP		Nov 16, 2027
		9555004		DP		Nov 16, 2027
		9622983		DP		Nov 16, 2027
		9622983		DP		Nov 16, 2027
<u>TOPIRAMATE - OUDEXY XR</u>						
N 205122	001	10363224				Mar 19, 2033
		8652527		DP		Mar 19, 2033
		8889190		DP		Mar 19, 2033
		9101545		DP		Mar 19, 2033
		9555005		DP		Mar 19, 2033
<u>TOPIRAMATE - OUDEXY XR</u>						
N 205122	002	10363224				Mar 19, 2033
		8652527		DP		Mar 19, 2033
		8889190		DP		Mar 19, 2033
		9101545		DP		Mar 19, 2033
		9555005		DP		Mar 19, 2033
<u>TOPIRAMATE - OUDEXY XR</u>						
N 205122	003	10363224				Mar 19, 2033
		8652527		DP		Mar 19, 2033
		8889190		DP		Mar 19, 2033
		9101545		DP		Mar 19, 2033
		9555005		DP		Mar 19, 2033
<u>TOPIRAMATE - OUDEXY XR</u>						
N 205122	004	10363224				Mar 19, 2033
		8652527		DP		Mar 19, 2033
		8889190		DP		Mar 19, 2033
		9101545		DP		Mar 19, 2033
		9555005		DP		Mar 19, 2033
<u>TOPIRAMATE - OUDEXY XR</u>						
N 205122	005	10363224				Mar 19, 2033
		8652527		DP		Mar 19, 2033
		8889190		DP		Mar 19, 2033
		9101545		DP		Mar 19, 2033
		9555005		DP		Mar 19, 2033

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<u>TOPOTECAN HYDROCHLORIDE - HYCAMTIN</u>						
N 020981	001 8158645	Dec 10, 2024	DP			
<u>TOPOTECAN HYDROCHLORIDE - HYCAMTIN</u>						
N 020981	002 8158645	Dec 10, 2024	DP			
<u>TRABECTEDIN - YONDELIS</u>						
N 207953	001 8895557	Jan 07, 2028	DP		M-232	Jun 29, 2021
	8895557*PED	Jul 07, 2028			NCE	Oct 23, 2020
					ODE-100	Oct 23, 2022
					PED	Apr 23, 2021
					PED	Dec 29, 2021
					PED	Apr 23, 2023
<u>TRAMADOL HYDROCHLORIDE - RYZOLT</u>						
N 021745	001 6607748	Jun 29, 2020	DP			
	7988998	Oct 27, 2023	DP			
<u>TRAMADOL HYDROCHLORIDE - RYZOLT</u>						
N 021745	002 6607748	Jun 29, 2020	DP			
	7988998	Oct 27, 2023	DP			
<u>TRAMADOL HYDROCHLORIDE - RYZOLT</u>						
N 021745	003 6607748	Jun 29, 2020	DP			
	7988998	Oct 27, 2023	DP			
<u>TRAMADOL HYDROCHLORIDE - CONZIP</u>						
N 022370	001 7858118	Apr 11, 2022	DP U-1104			
<u>TRAMADOL HYDROCHLORIDE - CONZIP</u>						
N 022370	002 7858118	Apr 11, 2022	DP U-1104			
<u>TRAMADOL HYDROCHLORIDE - CONZIP</u>						
N 022370	003 7858118	Apr 11, 2022	DP U-1104			
<u>TRAMETINIB DIMETHYL SULFOXIDE - MEKINIST</u>						
N 204114	001 7378423	May 29, 2027	DS DP		I-745	Jun 22, 2020
	8580304	Jan 28, 2032	DP		I-778	Apr 30, 2021
	8703781	Oct 15, 2030	DS DP U-1712		I-781	May 04, 2021
	8703781	Oct 15, 2030	DS DP U-2020		M-246	Oct 06, 2022
	8703781	Oct 15, 2030	DS DP U-2037		ODE-148	Jun 22, 2024
	8703781	Oct 15, 2030	DS DP U-2302		ODE-182	Apr 30, 2025
	8703781	Oct 15, 2030	DS DP U-2305		ODE-183	May 04, 2025
	8835443	Jun 10, 2025	U-1581		ODE-48	May 29, 2020
	8835443	Jun 10, 2025	U-1582		ODE-57	Jan 08, 2021
	8835443	Jun 10, 2025	U-2020			
	8835443	Jun 10, 2025	U-2037			
	8835443	Jun 10, 2025	U-2302			
	8835443	Jun 10, 2025	U-2305			
	8952018	Oct 15, 2030	U-2020			
	9155706	Jan 28, 2032	DP			
	9271941	Jan 28, 2032	DP			
	9399021	Jan 28, 2032	DP			
<u>TRAMETINIB DIMETHYL SULFOXIDE - MEKINIST</u>						
N 204114	002 7378423	May 29, 2027	DS DP		I-745	Jun 22, 2020
	8580304	Jan 28, 2032	DP		I-778	Apr 30, 2021
	8703781	Oct 15, 2030	DS DP U-1712		I-781	May 04, 2021
	8703781	Oct 15, 2030	DS DP U-2020		M-246	Oct 06, 2022
	8703781	Oct 15, 2030	DS DP U-2037		ODE-148	Jun 22, 2024
	8703781	Oct 15, 2030	DS DP U-2302		ODE-182	Apr 30, 2025
	8703781	Oct 15, 2030	DS DP U-2305		ODE-183	May 04, 2025
	8835443	Jun 10, 2025	U-1581		ODE-48	May 29, 2020
	8835443	Jun 10, 2025	U-1582		ODE-57	Jan 08, 2021
	8835443	Jun 10, 2025	U-2020			
	8835443	Jun 10, 2025	U-2037			
	8835443	Jun 10, 2025	U-2302			
	8835443	Jun 10, 2025	U-2305			
	8952018	Oct 15, 2030	U-2020			
	9155706	Jan 28, 2032	DP			
	9271941	Jan 28, 2032	DP			

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<u>TRAMETINIB DIMETHYL SULFOXIDE - MEKINIST</u>						
N 204114	002	9399021	Jan 28, 2032	DP		
<u>TRAMETINIB DIMETHYL SULFOXIDE - MEKINIST</u>						
N 204114	003	7378423	May 29, 2027	DS DP	I-745	Jun 22, 2020
		8580304	Jan 28, 2032	DP	I-778	Apr 30, 2021
		8703781	Oct 15, 2030	DS DP	I-781	May 04, 2021
		8703781	Oct 15, 2030	DS DP	U-1712	
		8703781	Oct 15, 2030	DS DP	U-2020	M-246 Oct 06, 2022
		8703781	Oct 15, 2030	DS DP	U-2037	ODE-148 Jun 22, 2024
		8703781	Oct 15, 2030	DS DP	U-2302	ODE-182 Apr 30, 2025
		8703781	Oct 15, 2030	DS DP	U-2305	ODE-183 May 04, 2025
		8835443	Jun 10, 2025	U-1581	ODE-48	May 29, 2020
		8835443	Jun 10, 2025	U-1582	ODE-57	Jan 08, 2021
		8835443	Jun 10, 2025	U-2020		
		8835443	Jun 10, 2025	U-2037		
		8835443	Jun 10, 2025	U-2302		
		8835443	Jun 10, 2025	U-2305		
		8952018	Oct 15, 2030	U-2020		
		9155706	Jan 28, 2032	DP		
		9271941	Jan 28, 2032	DP		
		9399021	Jan 28, 2032	DP		
<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>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	6607748	Jun 29, 2020	DP		
		7829120	Mar 27, 2027	DP	U-796	
		8133893	Mar 13, 2029	DS DP		
<u>TRAZODONE HYDROCHLORIDE - OLEPTRO</u>						
N 022411	002	6607748	Jun 29, 2020	DP		
		7829120	Mar 27, 2027	DP	U-796	
		8133893	Mar 13, 2029	DS DP		
<u>TREPROSTINIL - REMODULIN</u>						
N 021272	001	10076505	Dec 16, 2024	DP		
		7999007	Mar 29, 2029	DP	U-1437	
		8653137	Sep 05, 2028	U-1437		
		8658694	Sep 05, 2028	U-1437		
		9199908	May 24, 2024	U-1771		

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<u>TREPROSTINIL - REMODULIN</u>						
N 021272 001	9593066	Dec 15, 2028	DS			
	9604901	Dec 15, 2028	DS			
	9713599	Dec 16, 2024			U-2036	
<u>TREPROSTINIL - REMODULIN</u>						
N 021272 002	10076505	Dec 16, 2024	DP			
	7999007	Mar 29, 2029	DP		U-1437	
	8653137	Sep 05, 2028			U-1437	
	8658694	Sep 05, 2028			U-1437	
	9199908	May 24, 2024			U-1771	
	9593066	Dec 15, 2028	DS			
	9604901	Dec 15, 2028	DS			
	9713599	Dec 16, 2024			U-2036	
<u>TREPROSTINIL - REMODULIN</u>						
N 021272 003	10076505	Dec 16, 2024	DP			
	7999007	Mar 29, 2029	DP		U-1437	
	8653137	Sep 05, 2028			U-1437	
	8658694	Sep 05, 2028			U-1437	
	9199908	May 24, 2024			U-1771	
	9593066	Dec 15, 2028	DS			
	9604901	Dec 15, 2028	DS			
	9713599	Dec 16, 2024			U-2036	
<u>TREPROSTINIL - REMODULIN</u>						
N 021272 004	10076505	Dec 16, 2024	DP			
	7999007	Mar 29, 2029	DP		U-1437	
	8653137	Sep 05, 2028			U-1437	
	8658694	Sep 05, 2028			U-1437	
	9199908	May 24, 2024			U-1771	
	9593066	Dec 15, 2028	DS			
	9604901	Dec 15, 2028	DS			
	9713599	Dec 16, 2024			U-2036	
<u>TREPROSTINIL - TYVASO</u>						
N 022387 001	8497393	Dec 15, 2028	DS			
	9339507	Mar 10, 2028	DP			
	9358240	May 05, 2028			U-1849	
	9593066	Dec 15, 2028	DS			
	9604901	Dec 15, 2028	DS			
<u>TREPROSTINIL - REMODULIN</u>						
N 208276 001	10076505	Dec 16, 2024	DP			
	9593066	Dec 15, 2028	DS			
	9604901	Dec 15, 2028	DS			
	9713599	Dec 16, 2024			U-2036	
<u>TREPROSTINIL - REMODULIN</u>						
N 208276 002	10076505	Dec 16, 2024	DP			
	9593066	Dec 15, 2028	DS			
	9604901	Dec 15, 2028	DS			
	9713599	Dec 16, 2024			U-2036	
<u>TREPROSTINIL - REMODULIN</u>						
N 208276 003	10076505	Dec 16, 2024	DP			
	9593066	Dec 15, 2028	DS			
	9604901	Dec 15, 2028	DS			
	9713599	Dec 16, 2024			U-2036	
<u>TREPROSTINIL - REMODULIN</u>						
N 208276 004	10076505	Dec 16, 2024	DP			
	9593066	Dec 15, 2028	DS			
	9604901	Dec 15, 2028	DS			
	9713599	Dec 16, 2024			U-2036	
<u>TREPROSTINIL DIOLAMINE - ORENITRAM</u>						
N 203496 001	7417070	Jul 30, 2026	DS		I-820	Oct 18, 2022
	7544713	Jul 14, 2024			ODE-272	Oct 18, 2026
	8252839	May 24, 2024	DP			
	8349892	Jan 22, 2031	DP			

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<u>TREPROSTINIL DIOLAMINE - ORENITRAM</u>						
N 203496 001	8410169	Feb 13, 2030	DP			
	8747897	Aug 11, 2031	DP U-2724			
	8747897	Aug 11, 2031	DP U-2725			
	9050311	May 24, 2024	DS DP			
	9278901	May 24, 2024		U-1475		
	9393203	Apr 27, 2026	DP U-1877			
	9422223	May 24, 2024	DP			
	9593066	Dec 15, 2028	DS			
	9604901	Dec 15, 2028	DS			
<u>TREPROSTINIL DIOLAMINE - ORENITRAM</u>						
N 203496 002	7417070	Jul 30, 2026	DS		I-820	Oct 18, 2022
	7544713	Jul 14, 2024		U-1475	ODE-272	Oct 18, 2026
	8252839	May 24, 2024	DP			
	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			
	9050311	May 24, 2024	DS DP			
	9278901	May 24, 2024		U-1475		
	9393203	Apr 27, 2026	DP U-1877			
	9422223	May 24, 2024	DP			
	9593066	Dec 15, 2028	DS			
	9604901	Dec 15, 2028	DS			
<u>TREPROSTINIL DIOLAMINE - ORENITRAM</u>						
N 203496 003	7417070	Jul 30, 2026	DS		I-820	Oct 18, 2022
	7544713	Jul 14, 2024		U-1475	ODE-272	Oct 18, 2026
	8252839	May 24, 2024	DP			
	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			
	9050311	May 24, 2024	DS DP			
	9278901	May 24, 2024		U-1475		
	9393203	Apr 27, 2026	DP U-1877			
	9422223	May 24, 2024	DP			
	9593066	Dec 15, 2028	DS			
	9604901	Dec 15, 2028	DS			
<u>TREPROSTINIL DIOLAMINE - ORENITRAM</u>						
N 203496 004	7417070	Jul 30, 2026	DS		I-820	Oct 18, 2022
	7544713	Jul 14, 2024		U-1475	ODE-272	Oct 18, 2026
	8252839	May 24, 2024	DP			
	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			
	9050311	May 24, 2024	DS DP			
	9278901	May 24, 2024		U-1475		
	9393203	Apr 27, 2026	DP U-1877			
	9422223	May 24, 2024	DP			
	9593066	Dec 15, 2028	DS			
	9604901	Dec 15, 2028	DS			
<u>TREPROSTINIL DIOLAMINE - ORENITRAM</u>						
N 203496 005	7417070	Jul 30, 2026	DS		I-820	Oct 18, 2022
	7544713	Jul 14, 2024		U-1475	ODE-272	Oct 18, 2026
	8252839	May 24, 2024	DP			
	8349892	Jan 22, 2031	DP			
	8410169	Feb 13, 2030	DP			
	8747897	Aug 11, 2031	DP U-2724			
	8747897	Aug 11, 2031	DP U-2725			
	9050311	May 24, 2024	DS DP			
	9278901	May 24, 2024		U-1475		
	9393203	Apr 27, 2026	DP U-1877			
	9422223	May 24, 2024	DP			

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<u>TREPROSTINIL DIOLAMINE - ORENITRAM</u>						
N 203496 005	9593066	Dec 15, 2028	DS			
	9604901	Dec 15, 2028	DS			
<u>TRETINOIN - RENOVA</u>						
N 021108 001	6531141	Mar 07, 2020				
<u>TRETINOIN - ALTRENO</u>						
N 209353 001	6517847	Aug 03, 2020	DP U-2368		NDF	Aug 23, 2021
<u>TRIAMCINOLONE ACETONIDE - TRIAMCINOLONE ACETONIDE</u>						
A 212384 001					CGT	May 30, 2020
<u>TRIAMCINOLONE ACETONIDE - TRIESENC</u>						
N 022048 001	6395294	Jan 13, 2020	DP U-846			
	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		NP	Oct 06, 2020
	9555048	Aug 04, 2031		U-2151		
<u>TRICLABENDAZOLE - EGATEN</u>						
N 208711 001					NCE ODE-228	Feb 13, 2024 Feb 13, 2026
<u>TRIFAROTENE - AKLIEF</u>						
N 211527 001	7807708	Oct 01, 2026	DS DP		NCE	Oct 04, 2024
	8227507	Dec 21, 2025		U-818		
	8470871	Dec 21, 2025		U-2639		
	9084778	May 30, 2033	DP U-134			
<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		NP ODE-149	Jun 29, 2020 Jun 29, 2024
<u>UBROGEPANT - UBRELVY</u>						
N 211765 001	10117836	Jan 30, 2035	DP		NCE	Dec 23, 2024
	8754096	Jul 19, 2032	DS DP U-2717			
	8912210	Nov 10, 2031	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		NCE	Dec 23, 2024
	8754096	Jul 19, 2032	DS DP U-2717			
	8912210	Nov 10, 2031	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		
	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			
<u>UMECLIDINIUM BROMIDE - INCRUSE ELLIPTA</u>						
N 205382 001	7488827	Dec 18, 2027	DS DP			
	7498440	Apr 27, 2025	DS DP			
	8113199	Oct 23, 2027	DP			
	8161968	Feb 05, 2028	DP			
	8183257	Jul 27, 2025		U-1476		
	8201556	Feb 05, 2029	DP			
	8309572	Apr 27, 2025		U-1476		
	8534281	Mar 08, 2030	DP			

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<u>UMECLIDINIUM BROMIDE - INCRUSE ELLIPTA</u>						
N 205382	001 8746242	Oct 11, 2030	DP			
	9333310	Oct 02, 2027	DP			
<u>UMECLIDINIUM BROMIDE; VILANTEROL TRIFENATATE - ANORO ELLIPTA</u>						
N 203975	001 7439393	May 21, 2025	DS DP U-1476		M-245	Jun 09, 2022
	7488827	Dec 18, 2027	DS DP			
	7498440	Apr 27, 2025	DS DP			
	7776895	Sep 11, 2022	DP			
	8113199	Oct 23, 2027	DP			
	8161968	Feb 05, 2028	DP			
	8183257	Jul 27, 2025		U-1476		
	8309572	Apr 27, 2025		U-1476		
	8511304	Jun 14, 2027	DP U-1476			
	8534281	Mar 08, 2030	DP			
	8746242	Oct 11, 2030	DP			
	9333310	Oct 02, 2027	DP			
	9750726	Nov 29, 2030	DP			
	RE44874	Mar 23, 2023	DS DP U-1476			
<u>UNOPROSTONE ISOPROPYL - RESCULA</u>						
N 021214	001 6458836	Jul 09, 2021		U-1315		
	6458836	Jul 09, 2021		U-333		
<u>UPADACITINIB - RINVOO</u>						
N 211675	001 10519164	Oct 17, 2036	DP		NCE	Aug 16, 2024
	8962629	Jan 15, 2031	DS U-2616			
	9951080	Oct 17, 2036	DS DP			
	9963459	Oct 17, 2036	DP			
	RE47221	Dec 01, 2030	DS			
<u>URIDINE TRIACETATE - VISTOGARD</u>						
N 208159	001 6258795	Jul 10, 2023	DP		NCE	Sep 04, 2020
	7776838	Aug 17, 2027		U-1791	ODE-104	Dec 11, 2022
<u>URIDINE TRIACETATE - XURIDEN</u>						
N 208169	001 6258795	Jul 10, 2023	DP		NCE	Sep 04, 2020
					ODE-98	Sep 04, 2022
<u>VALBENAZINE TOSYLATE - INGREGZA</u>						
N 209241	001 10065952	Oct 28, 2036	DS DP U-1995		NCE	Apr 11, 2022
	8039627	Oct 06, 2029	DS DP			
	8357697	Nov 08, 2027		U-1995		
<u>VALBENAZINE TOSYLATE - INGREGZA</u>						
N 209241	002 10065952	Oct 28, 2036	DS DP U-1995		NCE	Apr 11, 2022
	8039627	Oct 06, 2029	DS DP			
	8357697	Nov 08, 2027		U-1995		
<u>VALGANCICLOVIR HYDROCHLORIDE - VALCYTE</u>						
N 022257	001 8889109	Dec 11, 2027	DP			
	9642911	Dec 11, 2027	DP			
<u>VANCOMYCIN HYDROCHLORIDE - FIRVANO KIT</u>						
N 208910	001 10493028	Mar 13, 2035	DP			
<u>VANCOMYCIN HYDROCHLORIDE - FIRVANO KIT</u>						
N 208910	002 10493028	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			
<u>VANCOMYCIN HYDROCHLORIDE - VANCOMYCIN HYDROCHLORIDE</u>						
N 211962	002 10039804	Nov 06, 2035	DP U-282			
	10188697	Nov 06, 2035	DP U-282			
<u>VANCOMYCIN HYDROCHLORIDE - VANCOMYCIN HYDROCHLORIDE</u>						
N 211962	003 10039804	Nov 06, 2035	DP U-282			
	10188697	Nov 06, 2035	DP U-282			

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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	004	10039804	Nov 06, 2035	DP U-282		
		10188697	Nov 06, 2035	DP U-282		
<u>VANDETANIB - CAPRELSA</u>						
N 022405	001	7173038	Aug 14, 2021	DS DP		
		8067427	Aug 08, 2028	DP		
		8642608	Feb 06, 2022		U-1490	
		RE42353	Jun 27, 2022	DS DP		
<u>VANDETANIB - CAPRELSA</u>						
N 022405	002	7173038	Aug 14, 2021	DS DP		
		8067427	Aug 08, 2028	DP		
		8642608	Feb 06, 2022		U-1490	
		RE42353	Jun 27, 2022	DS DP		
<u>VARDENAFIL HYDROCHLORIDE - LEVITRA</u>						
N 021400	001	8273876	Jul 23, 2027		U-1288	
		8841446	Jul 03, 2023	DP		
<u>VARDENAFIL HYDROCHLORIDE - LEVITRA</u>						
N 021400	002	8273876	Jul 23, 2027		U-1288	
		8841446	Jul 03, 2023	DP		
<u>VARDENAFIL HYDROCHLORIDE - LEVITRA</u>						
N 021400	003	8273876	Jul 23, 2027		U-1288	
		8841446	Jul 03, 2023	DP		
<u>VARDENAFIL HYDROCHLORIDE - LEVITRA</u>						
N 021400	004	8273876	Jul 23, 2027		U-1288	
		8841446	Jul 03, 2023	DP		
<u>VARDENAFIL HYDROCHLORIDE - STAXYN</u>						
N 200179	001	8613950	Dec 23, 2028	DP		
<u>VARENICLINE TARTRATE - CHANTIX</u>						
N 021928	001	6410550	May 10, 2020	DS DP U-56	M-237	Feb 22, 2022
		6410550*PED	Nov 10, 2020		PED	Aug 22, 2022
		6890927	May 06, 2022	DS DP U-56		
		6890927*PED	Nov 06, 2022			
		7265119	Aug 03, 2022	DS DP U-56		
		7265119*PED	Feb 03, 2023			
<u>VARENICLINE TARTRATE - CHANTIX</u>						
N 021928	002	6410550	May 10, 2020	DS DP U-56	M-237	Feb 22, 2022
		6410550*PED	Nov 10, 2020		PED	Aug 22, 2022
		6890927	May 06, 2022	DS DP U-56		
		6890927*PED	Nov 06, 2022			
		7265119	Aug 03, 2022	DS DP U-56		
		7265119*PED	Feb 03, 2023			
<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>VEMURAFENIB - ZELBORAF</u>						
N 202429	001	7504509	Oct 22, 2026	DS DP	I-757	Nov 06, 2020
		7863288	Jun 20, 2029	DS DP	ODE-158	Nov 06, 2024
		8143271	Jun 21, 2026	DS DP		
		8470818	Aug 02, 2026		U-1418	

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<u>VEMURAFENIB - ZELBORAF</u>						
N 202429 001	8470818	Aug 02, 2026			U-2164	
	8741920	Jul 27, 2030	DS DP			
	9447089	Jun 06, 2032	DP			
<u>VENETOCLAX - VENCLEXTA</u>						
N 208573 001	8546399	Jun 27, 2031	DS DP		I-782	Jun 08, 2021
	8722657	Jan 29, 2032	DS		I-789	Nov 21, 2021
	9174982	May 26, 2030		U-2323	I-795	May 15, 2022
	9174982	May 26, 2030		U-2445	M-228	Jun 08, 2021
	9174982	May 26, 2030		U-2446	NCE	Apr 11, 2021
	9174982	May 26, 2030		U-2537	ODE-114	Apr 11, 2023
	9539251	Sep 06, 2033		U-2538	ODE-185	Jun 08, 2025
					ODE-211	Nov 21, 2025
					ODE-239	May 15, 2026
<u>VENETOCLAX - VENCLEXTA</u>						
N 208573 002	8546399	Jun 27, 2031	DS DP		I-782	Jun 08, 2021
	8722657	Jan 29, 2032	DS		I-789	Nov 21, 2021
	9174982	May 26, 2030		U-2323	I-795	May 15, 2022
	9174982	May 26, 2030		U-2445	M-228	Jun 08, 2021
	9174982	May 26, 2030		U-2446	NCE	Apr 11, 2021
	9174982	May 26, 2030		U-2537	ODE-114	Apr 11, 2023
	9539251	Sep 06, 2033		U-2538	ODE-185	Jun 08, 2025
					ODE-211	Nov 21, 2025
					ODE-239	May 15, 2026
<u>VENETOCLAX - VENCLEXTA</u>						
N 208573 003	8546399	Jun 27, 2031	DS DP		I-782	Jun 08, 2021
	8722657	Jan 29, 2032	DS		I-789	Nov 21, 2021
	9174982	May 26, 2030		U-2323	I-795	May 15, 2022
	9174982	May 26, 2030		U-2445	M-228	Jun 08, 2021
	9174982	May 26, 2030		U-2446	NCE	Apr 11, 2021
	9174982	May 26, 2030		U-2537	ODE-114	Apr 11, 2023
	9539251	Sep 06, 2033		U-2538	ODE-185	Jun 08, 2025
					ODE-211	Nov 21, 2025
					ODE-239	May 15, 2026
<u>VILAZODONE HYDROCHLORIDE - VIIBRYD</u>						
N 022567 001	7834020	Jun 05, 2022	DS DP	U-839	M-254	Jan 31, 2023
	7834020*PED	Dec 05, 2022			PED	Jul 31, 2023
	8193195	Jun 05, 2022		U-839		
	8193195*PED	Dec 05, 2022				
	8236804	Jun 05, 2022		U-839		
	8236804*PED	Dec 05, 2022				
	8673921	Jun 05, 2022	DS DP			
	8673921*PED	Dec 05, 2022				
<u>VILAZODONE HYDROCHLORIDE - VIIBRYD</u>						
N 022567 002	7834020	Jun 05, 2022	DS DP	U-839	M-254	Jan 31, 2023
	7834020*PED	Dec 05, 2022			PED	Jul 31, 2023
	8193195	Jun 05, 2022		U-839		
	8193195*PED	Dec 05, 2022				
	8236804	Jun 05, 2022		U-839		
	8236804*PED	Dec 05, 2022				
	8673921	Jun 05, 2022	DS DP			
	8673921*PED	Dec 05, 2022				
<u>VILAZODONE HYDROCHLORIDE - VIIBRYD</u>						
N 022567 003	7834020	Jun 05, 2022	DS DP	U-839	M-254	Jan 31, 2023
	7834020*PED	Dec 05, 2022			PED	Jul 31, 2023
	8193195	Jun 05, 2022		U-839		
	8193195*PED	Dec 05, 2022				
	8236804	Jun 05, 2022		U-839		
	8236804*PED	Dec 05, 2022				
	8673921	Jun 05, 2022	DS DP			
	8673921*PED	Dec 05, 2022				
<u>VINCRISTINE SULFATE - MARQIBO KIT</u>						
N 202497 001	6723338	Mar 31, 2020			U-1271	
	7247316	Sep 25, 2020	DP			

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<u>VINCRIStINE SULFATE - MARQIBO KIT</u>						
N 202497 001	7887836	Mar 31, 2020	U-1271			
<u>VISMODEGIB - ERIVEDGE</u>						
N 203388 001	7888364	Nov 11, 2028	DS DP			
	9278961	Dec 15, 2028	U-1825			
<u>VORAPAXAR SULFATE - ZONTIVITY</u>						
N 204886 001	7235567	Jun 13, 2021	DS DP			
	7304078	Apr 06, 2024	DS DP	U-1512		
	7713999	May 30, 2024	DS DP	U-2291		
<u>VORICONAZOLE - VFEND</u>						
N 021266 001					NPP	Jan 29, 2022
<u>VORICONAZOLE - VFEND</u>						
N 021266 002					NPP	Jan 29, 2022
<u>VORICONAZOLE - VFEND</u>						
N 021267 001					NPP	Jan 29, 2022
<u>VORICONAZOLE - VFEND</u>						
N 021630 001					NPP	Jan 29, 2022
<u>VORINOSTAT - ZOLINZA</u>						
N 021991 001	7399787	Feb 09, 2025	U-892			
	7456219	Mar 11, 2027	DS			
	7652069	Mar 04, 2023	DP			
	7732490	Mar 04, 2023	U-892			
	7851509	Feb 21, 2024	DP	U-892		
	8067472	Mar 04, 2023	U-892			
	8093295	May 16, 2026	DP			
	8101663	Mar 04, 2023	U-892			
	8450372	Mar 18, 2028	U-892			
<u>VORTIOXETINE HYDROBROMIDE - TRINTELLIX</u>						
N 204447 001	7144884	Jun 17, 2026	DS DP	U-1439	M-227	May 02, 2021
	8476279	Oct 02, 2022	DP	U-1439	M-234	Oct 19, 2021
	8722684	Jun 30, 2031	DS DP			
	8969355	Jun 15, 2027		U-1668		
	9125908	Jun 15, 2027		U-2309		
	9125909	Jun 15, 2027		U-2309		
	9125910	Jun 15, 2027		U-2309		
	9227946	Jun 15, 2027		U-1668		
	9278096	Mar 21, 2032		U-2436		
	9861630	Jun 15, 2027		U-1668		
<u>VORTIOXETINE HYDROBROMIDE - TRINTELLIX</u>						
N 204447 002	7144884	Jun 17, 2026	DS DP	U-1439	M-227	May 02, 2021
	8476279	Oct 02, 2022	DP	U-1439	M-234	Oct 19, 2021
	8722684	Jun 30, 2031	DS DP			
	8969355	Jun 15, 2027		U-1668		
	9125908	Jun 15, 2027		U-2309		
	9125909	Jun 15, 2027		U-2309		
	9125910	Jun 15, 2027		U-2309		
	9227946	Jun 15, 2027		U-1668		
	9278096	Mar 21, 2032		U-2436		
	9861630	Jun 15, 2027		U-1668		
<u>VORTIOXETINE HYDROBROMIDE - TRINTELLIX</u>						
N 204447 003	7144884	Jun 17, 2026	DS DP	U-1439	M-227	May 02, 2021
	8476279	Oct 02, 2022	DP	U-1439	M-234	Oct 19, 2021
	8722684	Jun 30, 2031	DS DP			
	8969355	Jun 15, 2027		U-1668		
	9125908	Jun 15, 2027		U-2309		
	9125909	Jun 15, 2027		U-2309		
	9125910	Jun 15, 2027		U-2309		
	9227946	Jun 15, 2027		U-1668		
	9278096	Mar 21, 2032		U-2436		
	9861630	Jun 15, 2027		U-1668		

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<u>VORTIOXETINE HYDROBROMIDE - TRINTELLIX</u>						
N 204447	004	7144884	Jun 17, 2026	DS DP U-1439	M-227	May 02, 2021
		8476279	Oct 02, 2022	DP U-1439	M-234	Oct 19, 2021
		8722684	Jun 30, 2031	DS DP		
		8969355	Jun 15, 2027	U-1668		
		9125908	Jun 15, 2027	U-2309		
		9125909	Jun 15, 2027	U-2309		
		9125910	Jun 15, 2027	U-2309		
		9227946	Jun 15, 2027	U-1668		
		9278096	Mar 21, 2032	U-2436		
		9861630	Jun 15, 2027	U-1668		
<u>VOXELOTOR - OXBRYTA</u>						
N 213137	001	10017491	Dec 28, 2032	DP	NCE	Nov 25, 2024
		10034879	Dec 28, 2032	DS DP	ODE-281	Nov 25, 2026
		10493035	Oct 12, 2037	DP		
		9018210	Dec 28, 2032	DS DP		
		9248199	Jan 29, 2034	U-2676		
		9248199	Jan 29, 2034	U-2715		
		9447071	Feb 06, 2035	DS DP		
<u>ZANUBRUTINIB - BRUKINSA</u>						
N 213217	001	9447106	Apr 22, 2034	DS DP U-2145	NCE	Nov 14, 2024
					ODE-276	Nov 14, 2026
<u>ZICONOTIDE ACETATE - PRIALT</u>						
N 021060	001	8653033	Oct 01, 2024	U-48		
		8653033	Oct 01, 2024	U-55		
		8765680	Oct 01, 2024	U-48		
		8765680	Oct 01, 2024	U-55		
		9707270	Oct 01, 2024	U-2084		
<u>ZICONOTIDE ACETATE - PRIALT</u>						
N 021060	002	8653033	Oct 01, 2024	U-48		
		8653033	Oct 01, 2024	U-55		
		8765680	Oct 01, 2024	U-48		
		8765680	Oct 01, 2024	U-55		
		9707270	Oct 01, 2024	U-2084		
<u>ZICONOTIDE ACETATE - PRIALT</u>						
N 021060	003	8653033	Oct 01, 2024	U-48		
		8653033	Oct 01, 2024	U-55		
		8765680	Oct 01, 2024	U-48		
		8765680	Oct 01, 2024	U-55		
		9707270	Oct 01, 2024	U-2084		
<u>ZICONOTIDE ACETATE - PRIALT</u>						
N 021060	004	8653033	Oct 01, 2024	U-48		
		8653033	Oct 01, 2024	U-55		
		8765680	Oct 01, 2024	U-48		
		8765680	Oct 01, 2024	U-55		
		9707270	Oct 01, 2024	U-2084		
<u>ZIPRASIDONE HYDROCHLORIDE - GEODON</u>						
N 021483	001	7175855	May 18, 2020	DP		
<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		
		8052987	Oct 27, 2023	U-1199		

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<u>ZOLMITRIPTAN - ZOMIG</u>						
N 021450	003	6750237				
		Nov 28, 2020	DP			
		6750237*PED				
		May 28, 2021				
		7220767				
		Nov 28, 2020	DP			
		7220767*PED				
		May 28, 2021				
<u>ZOLMITRIPTAN - ZOMIG</u>						
N 021450	004	6750237				
		Nov 28, 2020	DP			
		7220767				
		Nov 28, 2020	DP			
<u>ZOLPIDEM TARTRATE - EDLUAR</u>						
N 021997	001	9265720				
		Feb 25, 2031				U-674
		9597281				U-674
		Apr 06, 2027				
<u>ZOLPIDEM TARTRATE - EDLUAR</u>						
N 021997	002	9265720				
		Feb 25, 2031				U-674
		9597281				U-674
		Apr 06, 2027				
<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				U-1194
		Feb 16, 2025				
		8242131				U-1266
		Aug 20, 2029				
		8252809				DP
		Feb 16, 2025				
<u>ZOLPIDEM TARTRATE - INTERMEZZO</u>						
N 022328	002	7658945				
		Apr 15, 2027	DP			U-1194
		7682628				U-1194
		Feb 16, 2025				
		8242131				U-1266
		Aug 20, 2029				
		8252809				DP
		Feb 16, 2025				

PATENT AND EXCLUSIVITY TERMS

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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)
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
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/M ² OR 175MG/M ² INTRAVENOUSLY OVER THREE HOURS EVERY THREE WEEKS
D-25	ADDITIONAL DOSAGE REGIMEN EQUAL TO HALF THE ORIGINAL DOSING REGIMEN

PATENT AND EXCLUSIVITY TERMS

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EXCLUSIVITY DOSING SCHEDULE

- 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 THIOPIENTAL
- 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
- 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

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY DOSING SCHEDULE**

- 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
- 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

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EXCLUSIVITY DOSING SCHEDULE

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 ALLERIC 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 SULATE CO-ADMINISTERED WITH RITONAVIR FOR THE TREATMENT OF HIV-1 INFECTION IN TREATMENT NAIVE PATIENTS
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

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EXCLUSIVITY DOSING SCHEDULE

- 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.
- 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

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- 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
- 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

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EXCLUSIVITY DOSING SCHEDULE

- 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

EXCLUSIVITY INDICATION

- I-1 DYSMENORRHEA
- I-2 CHOLANGIOPANCREATOGRAPHY
- I-3 INTRAVENOUS DIGITAL SUBTRACTION ANGIOGRAPHY
- I-4 PERIPHERAL VENOGRAPHY (PHLEBOGRAPHY)
- I-5 HYSTEROSALPINGOGRAPHY
- 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 VEIN 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

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

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
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	EROSIVE 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

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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 EROSIVE 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

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

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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

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

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- 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
- 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

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- 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 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

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- 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)
- 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 REVASCLARIZATION, 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

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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

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

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- 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 ADMINISTERED 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 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

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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
- 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

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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 HISTOPLASMOIS

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

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

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- 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
- 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>=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 &

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- 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 ABSCESSSES, 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
- I-401 LONGER-TERM EFFICACY OF ARIPIRAZOLE 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<=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

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- 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 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 REVASCULARIZATION 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 EROSIVE 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

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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 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

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- 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
- 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.

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- 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
- 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

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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

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

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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 NONSQUAMOUS NON-SMALL CELL LUNG CANCER
- 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

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- 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 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

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EXCLUSIVITY INDICATION

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

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

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EXCLUSIVITY INDICATION

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
- 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

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EXCLUSIVITY INDICATION

- 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 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

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EXCLUSIVITY INDICATION

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.
- 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

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EXCLUSIVITY INDICATION

- 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
- 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

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- 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
- 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

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EXCLUSIVITY INDICATION

- 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
- 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

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EXCLUSIVITY INDICATION

- 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
- 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 >= 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 RIVAROXBAN 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)

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

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- 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
- 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

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- 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 REVISIONS 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
- 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

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- 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 WITH 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 INSPRA (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 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

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- 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 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-109 CHANGES TO THE PACKAGE INSERT TO REFLECT THE RESULTS OF THE STUDY OF HEART AND RENAL PROTECTION (SHARP) TRIAL
- M-110 CHANGES TO THE PACKAGE INSERT TO REFLECT THE RESULTS OF THE STUDY OF HEART AND RENAL PROTECTION (SHARP) TRIAL

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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 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

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- 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
- 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

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- 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 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

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- 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.
- 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

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- 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
- 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

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- THE EFFECT OF VALGANCYCLOVIR 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/EDOXYBAN 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 BRf117277, A NON-RANDOMIZED, OPEN-LABEL, MULTI-CENTER, MULTI-COHORT TRIAL OF DABRAFENIB PLUS TRAMETINIB IN SUBJECTS WITH BRAF MUTATION-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

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- 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

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- 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-18 ADJUNCTIVE TREATMENT OF SEIZURES ASSOCIATED WITH LENNOX-GASTAUT SYNDROME INPATIENTS 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 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.

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- 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.
- 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

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- 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
- 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

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- 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
- 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

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- 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
- 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

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- 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
- 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 PATIENTS 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 PATIENTS WITH METASTATIC

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- 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 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

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- 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
- 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

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- 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 PARAGANGLIOMA 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
- 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

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- 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 INDICATED FOR THE TREATMENT OF LAMBERT-EATON MYASTHENIC SYNDROME (LEMS) IN ADULTS
- 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 NONSMALL CELL LUNG CANCER WHOSE TUMORS HAVE NON-RESISTANT EPIDERMAL GROWTH FACTOR 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 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

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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
- 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

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- 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 \geq 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
- ODE-277 TX OF ADULTS W/ ADV OVARIAN, FALLOPIAN TUBE, OR PRIMARY PERITONEAL CANCER TREATED W/ \geq 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/ \geq 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

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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

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

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U-20 A PROCESS FOR TREATING A PATIENT SUFFERING FROM PARKINSON'S SYNDROME AND IN NEED OF TREATMENT

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

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U-25 REDUCING CHOLESTEROL IN CHOLELITHIASIS PATIENTS

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U-28 CEREBRAL, CORONARY, PERIPHERAL, VISCERAL AND RENAL ARTERIOGRAPHY, AORTOGRAPHY AND LEFT VENTRICULOGRAPHY

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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

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U-35 TREATING CYTOMEGALOVIRUS IN A HUMAN WITH AN INJECTABLE COMPOSITION

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U-38 TREATMENT OF PAROXYSMAL SUPRAVENTRICULAR TACHYCARDIA

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U-45 TREATMENT OF INFLAMMATION AND ANALGESIA

U-46 TREATMENT OF PANIC DISORDER

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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

U-61 CEREBRAL AND PERIPHERAL ARTERIOGRAPHY AND CT IMAGING OF THE HEAD

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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

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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 RETINOPATY

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

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

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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

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

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PATENT USE

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

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

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PATENT USE

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

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

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PATENT USE

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

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

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PATENT USE

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 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

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PATENT USE

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

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 CONDITONS EMPLOYING OLANZAPINE

U-328 METHOD OF TREATING A PATIENT SUFFERING FROM ANY OF A NUMBER OF LISTED CONDITIONS

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- 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
- 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

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PATENT USE

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

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

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PATENT USE

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 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

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PATENT USE

- 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
- 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

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PATENT USE

- 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
- U-478 METHOD OF TREATING HEPATITIS C VIRAL INFECTION BY CONTINUOUS PARENTERAL ADMIN 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

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PATENT USE

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

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

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PATENT USE

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 IMMUNOCOPETENT 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 RESONACE 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.

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

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PATENT USE

- 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 SEGMENT OF THE GLOBE.
- U-576 ALLEX 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

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- 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 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.

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PATENT USE

- 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
- 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

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- 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
- 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

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PATENT USE

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 DYSLIPOPTEINEMIA 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 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

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PATENT USE

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 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

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PATENT USE

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 SEQUESTANT 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 ARIPIRAZOLE 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
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

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PATENT USE

- 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 PERSISTANT 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
- 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

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PATENT USE

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

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

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PATENT USE

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
- 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 HYPONATREMIA

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PATENT USE

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

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

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PATENT USE

- 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)
- 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

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PATENT USE

- 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
- 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

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PATENT USE

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
- 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 MIXED 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

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PATENT USE

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

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

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PATENT USE

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

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

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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 TROSPIDIUM 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., SALIVARRHEA) 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

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

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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 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

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- 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 TRETMENT 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
- 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

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- 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
- 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

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- 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)
- 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

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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

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 SULFONYUREA

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

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PATENT USE

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 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

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PATENT USE

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

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

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PATENT USE

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 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

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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
- 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

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PATENT USE

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

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

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PATENT USE

- 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) 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

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PATENT USE

- 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 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

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PATENT USE

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 RYTHM 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).
- 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

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PATENT USE

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.
- 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

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PATENT USE

U-1528 A METHOD OF LOWERING INTRAOCULAR PRESSURE

U-1529 ADJUNCTIVE TREATMENT OF MAJOR DEPRESSIVE DISORDER (MDD)

U-1530 USE OF ARIPIRAZOLE 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

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.

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- 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 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

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PATENT USE

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 ELVATED 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 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

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PATENT USE

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 ARIPIPIRAZOLE 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 PARITAPREVR 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 PARITAPREVR, OMBITASVIR, RITONAVIR, AND DASABUVIR WITH RIBAVIRIN.

U-1638 TREATMENT OF HCV INFECTION USING PARITAPREVR

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

U-1644 TREATMENT OF OVERACTIVE BLADDER BY APPLICATION OF OXYBUTYNYN 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

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PATENT USE

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

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

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PATENT USE

- 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
- 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.

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PATENT USE

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 THEAPY

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

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PATENT USE

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

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

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PATENT USE

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

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

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PATENT USE

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 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 PARITAPREVRIR, OMBITASVIR, RITONAVIR, AND DASABUVIR, WITHOUT RIBAVIRIN

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PATENT USE

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

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

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PATENT USE

- 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
- 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.

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PATENT USE

- 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
- 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

PATENT AND EXCLUSIVITY TERMS

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PATENT USE

- 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 CY1A2 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)
- 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

PATENT AND EXCLUSIVITY TERMS

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PATENT USE

- 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
- 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

PATENT AND EXCLUSIVITY TERMS

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PATENT USE

- 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
- 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

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PATENT USE

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

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

PATENT AND EXCLUSIVITY TERMS

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PATENT USE

- 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 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

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PATENT USE

- 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
- 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

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PATENT USE

- 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
- 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

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PATENT USE

- 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 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

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PATENT USE

- 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
- 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

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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

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

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PATENT USE

- 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 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

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PATENT USE

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 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

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- 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 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

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- 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
- 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

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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) 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

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- 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
- 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

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- 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 MYELOYDYSPLASIA-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 MYELOYDYSPLASIA-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
- 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)

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- U-2358 TREATMENT OF PATIENTS WITH HORMONE RECEPTOR (HR)-NEGATIVE BREAST CANCER WITH DELETERIOUS OR SUSPECTED DELETERIOUS GBRCA, 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 INIBITS 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 GLOBOTRIAOSYLCERAMIDE (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 LUMACAFTOR AND IVACAFTOR
- 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 LUMACAFTOR FORM I AND IVACAFTOR
- 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 LUMACAFTOR AND A SOLID COMPOSITION COMPRISING AMORPHOUS AND LESS THAN ABOUT 30% CRYSTALLINE IVACAFTOR
- U-2377 USE OF VITAL DYE FOR FACILITATING SURGICAL PROCEDURES FOR VITREO-RETINAL SURGERY
- 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

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- 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
- 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

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- 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
- 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

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- 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
- 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

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- 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
- 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

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PATENT USE

- 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
- 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

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PATENT USE

- 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
- 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

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PATENT USE

- 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
- 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 CARBOXYMALTOSE 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 CARBOXYMALTOSE
- U-2557 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 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

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PATENT USE

- 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
- 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 XPOVIO IS INDICATED IN COMBINATION WITH DEXAMETHASONE TO TREAT RELAPSED OR REFRACTORY MULTIPLE MYELOMA (REFRACTORY TO AT LEAST AN ANTI-CD38 MAB, 2

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PATENT USE

- 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
- 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

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PATENT USE

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
- 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

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- 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% OXYBUTYNYNIN 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 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

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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

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

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- 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 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 CARBOXYMALTOSIDE
- U-2710 A METHOD OF TREATING IRON DEFICIENCY ANEMIA IN ADULT PATIENTS WHO HAVE NON-DIALYSIS DEPENDENT CHRONIC KIDNEY DISEASE BY INTRAVENOUSLY ADMINISTERING FERRIC CARBOXYMALTOSIDE

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- 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 CARBOXYMALTOSIDE 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 CARBOXYMALTOSIDE 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 (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 DERISOMALTOSIDE
- 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)

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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