Our STN: BL103676/5137

SUPPLEMENT APPROVAL
May 2, 2023

Pharmacia & Upjohn Company LLC (A Pfizer Company)
Attention: William Vogt
235 East 42nd Street
New York, NY 10017

Dear Mr. Vogt:

We have approved your request received October 31, 2022, to supplement your Biologics License Application (BLA) submitted under Section 351(a) of the Public Health Service Act for the ATGAM® Sterile Solution, Lymphocyte Immune Globulin, Anti-thymocyte Globulin (Equine) to revise its United States Prescribing Information (USPI) based on the submitted post-marketing safety data. The USPI revisions for the ATGAM® Sterile Solution, Lymphocyte Immune Globulin, Anti-thymocyte Globulin (Equine) include revisions made to the Boxed Warning, Section 2 (Dosage and Administration), Section 4 (Contraindications), Section 5 (Warnings and Precautions), Section 6 (Post-Marketing Experience), Section 7 (Drug Interactions), Section 8 (Use in Specific Populations), Section 13 (Nonclinical Toxicology), and Section 17 (Patient Counseling Information).

LABELING

We hereby approve the draft content of labeling: Package Insert submitted under amendment 5002, dated May 2, 2023.

CONTENT OF LABELING

As soon as possible, but no later than 14 days from the date of this letter, please submit the final content of labeling (21 CFR 601.14) in Structured Product Labeling (SPL) format via the FDA automated drug registration and listing system, (eLIST) as described at http://www.fda.gov/ForIndustry/DataStandards/StructuredProductLabeling/default.htm. Content of labeling must be identical to the Package Insert submitted on May 2, 2023. Information on submitting SPL files using eLIST may be found in the guidance for industry *SPL Standard for Content of Labeling Technical Qs and As* at http://www.fda.gov/downloads/Drugs/GuidanceComplianceRegulatoryInformation/Guidances/UCM072392.pdf.

The SPL will be accessible via publicly available labeling repositories. All final labeling should be submitted as Product Correspondence to this BLA, STN BL 103676/0 at the time of use and include implementation information on Form FDA 356h.

ADVERTISING AND PROMOTIONAL LABELING.

You may submit two draft copies of the proposed introductory advertising and promotional labeling with Form FDA 2253 to the Advertising and Promotional Labeling Branch at the following address:

Food and Drug Administration
Center for Biologics Evaluation and Research
Document Control Center
10903 New Hampshire Ave.
WO71–G112
Silver Spring, MD 20993-0002

You must submit copies of your final advertising and promotional labeling at the time of initial dissemination or publication, accompanied by Form FDA 2253 (21 CFR 601.12(f)(4)).

All promotional claims must be consistent with and not contrary to approved labeling. You should not make a comparative promotional claim or claim of superiority over other products unless you have substantial evidence or substantial clinical experience to support such claims (21 CFR 202.1(e)(6)).

Please submit an amendment to all pending supplemental applications for this BLA that include revised labeling incorporating a revised content of labeling that includes these changes.

We will include information contained in the above-referenced supplement in your BLA file.

Sincerely,

Celia Witten, MD
Acting Director
Division of Clinical Evaluation General Medicine
Office of Clinical Evaluation
Office of Therapeutic Products
Center for Biologics Evaluation and Research