Our STN: BL 103869/5738

SUPPLEMENT APPROVAL
January 27, 2026

Genzyme Corporation
Attention: Gabriel Joseph
50 Binney Street
Cambridge, MA 02141

Dear Gabriel Joseph:

We have approved your request received July 31, 2025, to supplement your Biologics License Application (BLA) submitted under section 351(a) of the Public Health Service Act for anti-thymocyte globulin (rabbit). The supplement revises the USPI as follows: Boxed Warning removed; Section 1 (Indications and Usage) specifies pediatric and adult populations; Section 5 (Warnings and Precautions) streamlines risk and mitigation information; Section 6.2 (Postmarketing Experience) adds anemia including hemolytic anemia, thrombotic microangiopathy, hepatic failure, and hyperbilirubinemia; and Section 8.4 (Pediatric Use) specifies supporting data for pediatric use of the USPI for Thymoglobulin®.

LABELING

We hereby approve the draft content of labeling Patient Package Insert submitted in your amendment received, January 22, 2026.

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 https://www.fda.gov/industry/fda-data-standards-advisory-board/structured-product-labeling-resources. Content of labeling must be identical to the Package Insert submitted on January 22, 2026. 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 https://www.fda.gov/regulatory-information/search-fda-guidance-documents/spl-standard-content-labeling-technical-qs.

The SPL will be accessible via publicly available labeling repositories.

All final labeling should be submitted as Product Correspondence to this BLA, STN BL 103869 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)).

For each pending supplemental application for this BLA that includes proposed revised labeling, please submit an amendment to update the proposed revised labeling with the changes approved today.

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

Sincerely,

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