Our STN: BL 125078/1003

SUPPLEMENT APPROVAL
January 30, 2024

CSL Behring LLC
Attention: Edward T. Gallagher
1201 North Kinzie Avenue
Bradley, IL 60915

Dear Edward Gallagher:

We have approved your request, received September 29, 2023, to supplement your Biologics License Application (BLA) submitted under section 351(a) of the Public Health Service Act for Alpha-1-Proteinase Inhibitor (Human) [ZEMAIRA], to update the specification limit for the chloride excipient from (b) (4) 30 - 39 mM for ZEMAIRA drug product manufactured and packaged primarily at CSL Behring LLC, Bradley, IL, with secondary labeling and packaging done at (b) (4)

LABELING

We hereby approve the draft content of labeling Package Insert submitted under amendment 0, dated September 29, 2023, and the draft carton and container labels submitted under amendment 2, dated January 23, 2024.

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 September 29, 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.

CARTON AND CONTAINER LABELS

Please electronically submit final printed carton and container labels identical to the carton and container labels submitted on January 23, 2024, according to the guidance

for industry *Providing Regulatory Submissions in Electronic Format — Certain Human Pharmaceutical Product Applications and Related Submissions Using the eCTD Specifications* at https://www.fda.gov/regulatory-information/search-fda-guidance-documents/providing-regulatory-submissions-electronic-format-certain-human-pharmaceutical-product-applications.

All final labeling should be submitted as Product Correspondence to this BLA, STN BL 125078 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,

Dorothy Scott, MD
Director
Division of Plasma Derivatives
Office of Plasma Protein Therapeutics
Office of Therapeutic Products
Center for Biologics Evaluation and Research