



BLA 761143/S-031

## SUPPLEMENT APPROVAL

Horizon Therapeutics Ireland DAC  
c/o Horizon Therapeutics USA Inc  
Attention: Nicole Cheung, PharmD  
Manager, Regulatory Affairs  
One Horizon Way  
Thousand Oaks, CA 91320

Dear Dr. Cheung:

Please refer to your supplemental biologics license application (sBLA) received July 29, 2025, and your amendments, submitted under section 351(a) of the Public Health Service Act for TEPEZZA (teprotumumab-trbw) for injection.

This Prior Approval supplemental biologics application provides for updates in the WARNINGS AND PRECAUTIONS and ADVERSE REACTIONS sections of the Prescribing Information to add postapproval reports of exacerbation of inflammatory bowel disease (IBD), including bowel perforation, in some patients without a prior diagnosis of IBD.

### **APPROVAL & LABELING**

We have completed our review of this application, as amended. It is approved, effective on the date of this letter, for use as recommended in the enclosed agreed-upon labeling.

### **CONTENT OF LABELING**

As soon as possible, but no later than 14 days from the date of this letter, submit, via the FDA automated drug registration and listing system (eLIST), the content of labeling [21 CFR 601.14(b)] in structured product labeling (SPL) format, as described at FDA.gov,<sup>1</sup> that is identical to the enclosed labeling (text for the Prescribing Information) and include the labeling changes proposed in any pending "Changes Being Effected" (CBE) supplements.

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*.<sup>2</sup>

The SPL will be accessible via publicly available labeling repositories.

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<sup>1</sup> <http://www.fda.gov/ForIndustry/DataStandards/StructuredProductLabeling/default.htm>

<sup>2</sup> We update guidances periodically. For the most recent version of a guidance, check the FDA Guidance Documents Database <https://www.fda.gov/RegulatoryInformation/Guidances/default.htm>.

Also within 14 days, amend all pending supplemental applications that include labeling changes for this BLA, including pending "Changes Being Effected" (CBE) supplements, for which FDA has not yet issued an action letter, with the content of labeling [21 CFR 601.12(f)] in Microsoft Word format that includes the changes approved in this supplemental application, as well as annual reportable changes. To facilitate review of your submission(s), provide a highlighted or marked-up copy that shows all changes, as well as a clean Microsoft Word version. The marked-up copy should provide appropriate annotations, including supplement number(s) and annual report date(s).

### **REPORTING REQUIREMENTS**

We remind you that you must comply with reporting requirements for an approved BLA (in 21 CFR 600.80 and in 21 CFR 600.81).

If you have any questions, contact Ahmed Ayodeji, PharmD, Regulatory Project Manager, at (301) 837-7390 or [Ahmed.Ayodeji@fda.hhs.gov](mailto:Ahmed.Ayodeji@fda.hhs.gov).

Sincerely,

*{See appended electronic signature page}*

William M. Boyd, MD  
Deputy Director  
Division of Ophthalmology  
Office of Specialty Medicine  
Office of New Drugs  
Center for Drug Evaluation and Research

### ENCLOSURE:

- Content of Labeling
  - Prescribing Information

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**This is a representation of an electronic record that was signed electronically. Following this are manifestations of any and all electronic signatures for this electronic record.**  
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/s/  
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WILLIAM M BOYD  
08/26/2025 03:33:43 PM