



NDA 211673/S-001, 211672/S-001

**SUPPLEMENT APPROVAL**

Nabriva Therapeutics Ireland DAC  
c/o Nabriva Therapeutics US, INC.  
Attention: Mersedeh Miraliakbari, PharmD  
Vice President, Regulatory Affairs  
1000 Continental Drive, Suite 600  
King of Prussia, PA 19406

Dear Dr. Miraliakbari:

Please refer to your supplemental New Drug Application(s) (sNDA) submitted under section 505(b) of the Federal Food, Drug, and Cosmetic Act (FDCA), and all amendments, for the following products:

<b>Supplemental Application</b>	<b>Product Information</b>	<b>Submit Date</b>	<b>FDA Received Date</b>
NDA 211673/S-001	Xenleta (lefamulin) Injection, 150 mg	April 15, 2020	April 15, 2020
NDA 211672/S-001	XENLETA (lefamulin) Tablets, 600 mg	April 15, 2020	April 15, 2020

These “Changes Being Effected” supplemental new drug applications provide for the following changes:

1. Change HOW SUPPLIED storage statement for the citrate buffer diluent bags to state store in overwrap at 2°C to 25°C (36°F to 77°F) until ready to use.
2. Add Tradename to diluent container label and carton.

**APPROVAL & LABELING**

We have completed our review of these supplemental applications, as amended. They are 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 the content of labeling [21 CFR 314.50(l)] in structured product labeling (SPL) format using 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 enclosed labeling (text for the prescribing

information) with the addition of any labeling changes in pending “Changes Being Effected” (CBE) supplements, as well as annual reportable changes not included in the enclosed labeling.

Information on submitting SPL files using eLIST may be found in the guidance for industry titled *SPL Standard for Content of Labeling Technical Qs and As* at <http://www.fda.gov/downloads/DrugsGuidanceComplianceRegulatoryInformation/Guidances/UCM072392.pdf>.

Information on submitting SPL files using eLIST may be found in the guidance for industry titled *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.

Also within 14 days, amend all pending supplemental applications that include labeling changes for this NDA, including CBE supplements for which FDA has not yet issued an action letter, with the content of labeling [21 CFR 314.50(I)(1)(i)] in MS Word format, that includes the changes approved in these supplemental applications, as well as annual reportable changes, and annotate each change. To facilitate review of your submission, 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).

### **CARTON AND CONTAINER LABELS**

Submit final printed carton and container labels that are identical to enclosed carton and container labels, as soon as they are available, but no more than 30 days after they are printed. Please submit these labels electronically according to the guidance for industry *Providing Regulatory Submissions in Electronic Format – Certain Human Pharmaceutical Product Applications and Related Submissions Using the eCTD Specifications*. For administrative purposes, designate this submission “**Product Correspondence – Final Printed Carton and Container Labels for approved NDA 211673/S-001.**” Approval of this submission by FDA is not required before the labeling is used.

**REPORTING REQUIREMENTS**

We remind you that you must comply with reporting requirements for an approved NDA (21 CFR 314.80 and 314.81).

If you have any questions, call Chinedu Ebonine, Regulatory Business Process Manager, at (240) 402 - 3448.

Sincerely,

*{See appended electronic signature page}*

David Lewis, Ph.D.  
Branch Chief, BII  
Division of Post-Marketing Activities I  
Office of Lifecycle Drug Products  
Office of Pharmaceutical Quality  
Center for Drug Evaluation and Research

Enclosure(s):

Content of Labeling  
Carton and Container Labeling



David  
Lewis

Digitally signed by David Lewis

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