



NDA 020388/S-37

## SUPPLEMENT APPROVAL

Pierre Fabre Medicament  
Attention: Kristen Snaguski  
Director of Quality Assurance  
Regulatory Affairs, Vigilance & Information  
8 Campus Dr, Suite 202  
Parsippany, NJ 07054

Dear Ms. Snaguski:

Please refer to your supplemental new drug application (sNDA) dated and received December 2, 2019, and your amendments dated January 15, 2020 and January 24, 2020, submitted under section 505(b) of the Federal Food, Drug, and Cosmetic Act (FDCA) for Navelbine (vinorelbine) injection.

This Prior Approval supplemental new drug application provides for the removal of instructions involving preparation of vinorelbine in a syringe from the Dosage and Administration subsection 2.3 of the package insert.

### **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 with minor editorial revisions listed below and reflected in the enclosed labeling.

- Highlights of Prescribing Information revised to include the date of Prescribing Information last updated as "1/2020".
- Removed extra space in the Dosage and Administration subsection 2.3 Preparation and Administration between the subheading "Preparation" and the next sentence "Dilute NAVELBINE in an intravenous bag to a concentration between 0.5 mg/mL and 2 mg/mL".

## **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 FDA.gov.<sup>1</sup> 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 Effectuated” (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 *SPL Standard for Content of Labeling Technical Qs and As*.<sup>2</sup>

The SPL will be accessible from 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(l)(1)(i)] 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).

## **REQUIRED PEDIATRIC ASSESSMENTS**

Under the Pediatric Research Equity Act (PREA) (21 U.S.C. 355c), all applications for new active ingredients (which includes new salts and new fixed combinations), new indications, new dosage forms, new dosing regimens, or new routes of administration are required to contain an assessment of the safety and effectiveness of the product for the claimed indication in pediatric patients unless this requirement is waived, deferred, or inapplicable.

Because none of these criteria apply to your application, you are exempt from this requirement.

## **REPORTING REQUIREMENTS**

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

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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>.

If you have any questions you may contact Felicia Diggs, Safety Regulatory Project Manager, at (240) 402-4932 or via email at Felicia.diggs@fda.hhs.gov.

Sincerely,

*{See appended electronic signature page}*

Shanthi Marur, M.B.B.S., M.D.  
Associate Director for Safety (acting)  
Office of Oncologic Diseases  
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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SHANTHI N MARUR  
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