



NDA 21632/S-024

SUPPLEMENT APPROVAL

Vicuron Holdings LLC, a subsidiary of Pfizer, Inc.
Attention: Nadia D. Kirzecky
Director, Pfizer Essential Health Global Regulatory Brands
235 East 42nd Street
New York, NY 10017

Dear Ms. Kirzecky:

Please refer to your Supplemental New Drug Application (sNDA) dated December 13, 2016, received December 13, 2016, and your amendments, submitted under section 505(b) of the Federal Food, Drug, and Cosmetic Act (FDCA) for ERAXIS (anidulafungin IV) 50 mg and 100 mg.

This Prior Approval supplemental new drug application provides for revisions to update the ERAXIS prescribing information (PI) to be in compliance with the FDA Pregnancy and Lactation Labeling Rule (PLLR). Specifically, revisions were made to the **USE IN SPECIFIC POPULATIONS (8)** section, Pregnancy (8.1) and Lactation (8.2), subsections; **PATIENT COUNSELING INFORMATION (17)** section, Pregnant Women and Breastfeeding Mothers (17.3) subsection. Additionally, editorial revisions were made to the **HIGHLIGHTS, TABLE OF CONTENTS, CLINICAL PHARMACOLOGY (12)** section, Pharmacokinetics (12.3) subsection, as well as minor editorial changes throughout the PI.

APPROVAL & LABELING

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

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 package insert), 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>.

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 MS Word format, that includes the changes approved in this supplemental application, 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).

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 Mr. Gregory DiBernardo, Regulatory Project Manager, at (301) 796-4063.

Sincerely,

{See appended electronic signature page}

Dmitri Iarikov, M.D., Ph.D.
Acting Deputy Director
Division of Anti-Infective Products
Office of Antimicrobial Products
Center for Drug Evaluation and Research

ENCLOSURE:
Content of Labeling

This is a representation of an electronic record that was signed electronically and this page is the manifestation of the electronic signature.

/s/

DMITRI IARIKOV
08/17/2017