

NDA 050441/S-086
NDA 050639/S-043

SUPPLEMENT APPROVALS

Pfizer, Inc.
Attention: Mikhail Abarshalin
Director, Pfizer Global Regulatory Affairs
235 East 42nd Street
New York, NY 10017-7555

Dear Mr. Abarshalin:

Please refer to your supplemental new drug applications (sNDA) dated and received October 14, 2021, submitted under section 505(b) of the Federal Food, Drug, and Cosmetic Act (FDCA) for the following:

- Cleocin Phosphate Sterile Solution, 150 mg/mL [NDA 050441]
- Cleocin Phosphate IV Sterile Solution, 300 mg/50 mL, 600 mg/50 mL and 900 mg/50mL [NDA 050639]

These "Changes Being Effected" sNDAs provide for revisions to the prescribing information in response to the Agency supplement request letter dated September 14, 2021, requiring the Pfizer, under Section 409I of the Public Health Service Act (also known as the Best Pharmaceuticals for Children Act (BPCA), which mandates the critical need for pediatric labeling) to update the label with pediatric information.

APPROVAL & LABELING

We have completed our review of these applications and 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(I)] in structured product labeling (SPL) format using the FDA automated drug registration and listing system (eLIST), as described at FDA.gov.¹

¹ <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 *SPL Standard for Content of Labeling Technical Qs and As*.²

The SPL will be accessible from publicly available labeling repositories.

Also within 14 days, amend all pending supplemental applications that include labeling changes for these NDAs, 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 these supplemental applications, 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 NDA (21 CFR 314.80 and 314.81).

If you have any questions, call J. Christopher Davi, MS, Senior Regulatory Project Manager, at (301) 796-0702.

Sincerely,

{See appended electronic signature page}

Dmitri Iarikov, MD, PhD
Deputy Director
Division of Anti-Infectives
Office of Infectious Diseases
Center for Drug Evaluation and Research

ENCLOSURE: Prescribing Information

² 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>.

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.

/s/

DMITRI IARIKOV
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