



NDA 50-558/S-075

SUPPLEMENT APPROVAL

Teligent OÜ
c/o Teligent Pharma, Inc.
Attention: Susan Todd, Director, Regulatory Affairs
105 Lincoln Avenue
P.O. Box 687
Buena, NJ 08310

Dear Ms. Todd:

Please refer to your supplemental new drug application (sNDA) dated March 22, 2019, received March 22, 2019, submitted under section 505(b) of the Federal Food, Drug, and Cosmetic Act (FDCA) for Zinacef (cefuroxime sodium) Intravenous, 0.75 g, 1.5 g, and 7.5 g.

This Prior Approval supplemental new drug application, submitted in response to our February 26, 2019, supplement request letter, provides for removal of the **Drug Interactions** subsection, from the **PRECAUTIONS** section, as data from clinical and pharmacokinetic studies do not provide evidence of drug interactions between hormonal contraceptives and non-rifamycin antibacterial drugs, including cephalosporins.

APPROVAL & LABELING

We have completed our review of this application and 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 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.¹ Content of labeling must be identical to the enclosed labeling, 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.

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

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 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).

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: Content of Labeling

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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