ANDA 215840

ANDA TENTATIVE APPROVAL

Eugia US LLC
U.S. Agent for Eugia Pharma Specialities Limited
279 Princeton-Hightstown Road
East Windsor, NJ 08520
Attention: Apexa Chudasama
Senior Director, Regulatory Affairs

Dear Apexa Chudasama:

This letter is in reference to your abbreviated new drug application (ANDA) received for review on February 8, 2021, submitted pursuant to section 505(j) of the Federal Food, Drug, and Cosmetic Act (FD&C Act) for Etelcalcetide Injection, 2.5 mg/0.5 mL, 5 mg/mL, and 10 mg/2 mL (5 mg/mL) Single-Dose Vials.

Reference is also made to the complete response letter issued by this office on December 1, 2022, and to any amendments thereafter.

We have completed the review of this ANDA and have concluded that adequate information has been presented to demonstrate that the drug meets the requirements for approval under the FD&C Act. We have determined your Etelcalcetide Injection, 2.5 mg/0.5 mL, 5 mg/mL, and 10 mg/2 mL (5 mg/mL) Single-Dose Vials to be bioequivalent and therapeutically equivalent to the reference listed drug (RLD), Parsabiv Injection, 2.5 mg/0.5 mL, 5 mg/mL, and 10 mg/2 mL (5 mg/mL), of KAI Pharmaceuticals, Inc. (KAI).

However, we are unable to grant final approval to your ANDA at this time because of the patent issue noted below. Therefore, the ANDA is **tentatively approved**. This determination is based upon information available to the Agency at this time (e.g., information in your ANDA and the status of current good manufacturing practices (cGMPs) of the facilities used in the manufacturing and testing of the drug product). This determination is subject to change on the basis of new information that may come to our attention. This letter does not address issues related to the 180-day exclusivity provisions under section 505(j)(5)(B)(iv) of the FD&C Act.

The reference listed drug (RLD) upon which you have based your ANDA, KAI's Parsabiv Injection, 2.5 mg/0.5 mL, 5 mg/mL, and 10 mg/2 mL (5 mg/mL), is subject to periods of patent protection. The following patents and expiration dates are currently listed in the Agency's publication titled *Approved Drug Products with Therapeutic Equivalence Evaluations* (the "Orange Book"):

U.S. Patent Number	Expiration Date
8,377,880 (the '880 patent)	July 29, 2030
8,999,932 (the '932 patent)	February 7, 2031
9,278,995 (the '995 patent)	July 29, 2030
9,701,712 (the '712 patent)	July 29, 2030
9,820,938 (the '938 patent)	June 27, 2034
10,344,765 (the '765 patent)	June 27, 2034
11,162,500 (the '500 patent)	June 27, 2034

With respect to the '880, '932, '995 and '712 patents, your ANDA contains paragraph III certifications to each of the patents under section 505(j)(2)(A)(vii)(III) of the FD&C Act stating that Eugia Pharma Specialities Limited (Eugia) will not market Etelcalcetide Injection, 2.5 mg/0.5 mL, 5 mg/mL, and 10 mg/2 mL (5 mg/mL) Single-Dose Vials prior to the expiration of the patents. Therefore, final approval of your ANDA may not be granted pursuant to section 505(j)(5)(B)(ii) of the FD&C Act until the '932 patent has expired, currently February 7, 2031.

Your ANDA contains paragraph IV certifications to the '938, '765 and '500 patents[1] under section 505(j)(2)(A)(vii)(IV) of the FD&C Act stating that the patents are invalid, unenforceable, or will not be infringed by your manufacture, use, or sale of Etelcalcetide Injection, 2.5 mg/0.5 mL, 5 mg/mL, and 10 mg/2 mL (5 mg/mL) Single-Dose Vials, under this ANDA. You have notified the Agency that Eugia complied with the requirements of section 505(j)(2)(B) of the FD&C Act. Litigation was initiated within the statutory 45-day period against Eugia for infringement of the '938 and '765 patents in the United States District Court for the District of Delaware [Amgen Inc. and KAI Pharmaceuticals, Inc. v. Aurobindo Pharma Limited, Aurobindo Pharma USA Inc., Eugia Pharma Specialties Limited, and AuroMedics Pharma LLC, Civil Action No. 21-00662].

Therefore, final approval cannot be granted until:

1. a. the expiration of the 7.5-year period provided for in sections 505(j)(5)(B)(iii) and 505(j)(5)(F)(ii) of the FD&C Act,

   b. the date the court decides[2] that the '938 and '765 'patents are invalid or not infringed (see sections 505(j)(5)(B)(iii)(I), (II), and (III) of the FD&C Act), or

   c. The '938, '765, '880, '932, '995, '712 and '500 patents have expired, and

2. The Agency is assured there is no new information that would affect whether final approval should be granted.

Please note that if FDA requires a Risk Evaluation and Mitigation Strategy (REMS) for a listed drug, an ANDA referencing that listed drug also will be required to have a REMS. See section 505-1(i) of the FD&C Act.

REQUIREMENTS AND RECOMMENDATIONS POST APPROVAL

Under applicable statutes, regulations, and guidances, if your ANDA receives final approval, it may be subject to certain requirements and recommendations post approval, including requirements regarding changes to approved ANDAs, postmarketing reporting, promotional materials, and annual facility fees, among others. For information on post-approval requirements and recommendations for ANDAs and a list of resources for ANDA holders, we refer you to https://www.fda.gov/drugs/abbreviated-new-drug-application-anda/requirements-and-resources-approved-andas.

RESUBMISSION

To request final approval, please submit an amendment titled "FINAL APPROVAL REQUESTED" with enough time to permit FDA review prior to the date you believe that your ANDA will be eligible for final approval. A request for final approval that contains no new data, information, or other changes to the ANDA generally requires a period of 3 months for Agency review. Accordingly, such a request for final approval should be submitted no later than 3 months prior to the date on which you seek approval. A request for final approval that contains substantive changes to this ANDA or changes in the status of the manufacturing and testing facilities' compliance with cGMPs will be classified and reviewed according to OGD policy in effect at the time of receipt. Applicants should review available agency guidance for industry related to amendments under the generic drug user fee program to determine the duration of Agency review needed to review the changes submitted. As part of this consideration, applicants should monitor any changes to the RLD that occur after tentative approval, including changes in labeling, patent or exclusivity information, or marketing status. The submission of multiple amendments prior to final approval may also result in a delay in the issuance of the final approval letter.

The amendment requesting final approval should provide the legal/regulatory basis for your request for final approval and should include a copy of a court decision, settlement or licensing agreement, or other information described in 21 CFR 314.107, as appropriate. It should also identify changes, if any, in the conditions under which the ANDA was tentatively approved, e.g., updated information such as final-printed labeling, chemistry, manufacturing, and controls data as appropriate. This amendment should be submitted even if none of these changes were made, and it should be designated clearly in your cover letter as a "FINAL APPROVAL REQUESTED."

In addition to the amendment requested above, the Agency may request, at any time prior to the date of final approval, that you submit an additional amendment containing information as specified by the Agency. Failure to submit either or, if requested, both types of amendments described above may result in a delay in the issuance of the final approval letter.

This drug product may not be marketed without final Agency approval under section 505(j) of the FD&C Act. The introduction or delivery for introduction into interstate commerce of this drug product before the final approval date is prohibited under section 301 of the FD&C Act. Also, until the Agency issues the final approval letter, this drug product will not be deemed approved for marketing under section 505(j) of the FD&C Act, and will not be listed in the Orange Book.

For further information on the status of this ANDA or upon submitting an amendment to the ANDA, please contact Devon Lee, Regulatory Project Manager, at (301) 837 - 7615.

Sincerely yours,

{See appended electronic signature page}

For Edward M. Sherwood
Director
Office of Regulatory Operations
Office of Generic Drugs
Center for Drug Evaluation and Research

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[1] The Agency notes that the '500 patent was submitted to the Agency after submission of your ANDA. Litigation, if any, with respect to this patent would not create a statutory stay of approval.

[2] This decision may be either a decision of the district court or the court of appeals, whichever court is the first to decide that the patent is invalid or not infringed.

Catherine Poole

Digitally signed by Catherine Poole
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