



ANDA 213154

ANDA TENTATIVE APPROVAL

Lupin Pharmaceuticals, Inc.
U.S. Agent for Natco Pharma Limited
111 South Calvert Street
Harborplace Tower, 21st Floor
Baltimore, MD 21202
Attention: Debashis Mohanty
Senior Manager - Regulatory Affairs

Dear Sir:

This letter is in reference to your abbreviated new drug application (ANDA) received for review on February 8, 2019, submitted pursuant to section 505(j) of the Federal Food, Drug, and Cosmetic Act (FD&C Act) for Bosentan Tablets for Oral Suspension, 32 mg.

Reference is also made to the complete response letter issued by this office on December 2, 2019, 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 Bosentan Tablets for Oral Suspension, 32 mg, to be bioequivalent and therapeutically equivalent to the reference listed drug (RLD), Tracleer Tablets for Oral Suspension, 32 mg, of Actelion Pharmaceuticals US, Inc. (Actelion).

However, we are unable to grant final approval to your ANDA at this time because of the patent and exclusivity issues 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 RLD upon which you have based your ANDA, Actelion's Tracleer Tablets for Oral Suspension, 32 mg, 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>U.S. Patent Number</u>	<u>Expiration Date</u>
7,959,945 (the '945 patent)	December 28, 2027
8,309,126 (the '126 patent)	May 15, 2026

Your ANDA contains paragraph IV certifications to each of the patents 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 Bosentan Tablets for Oral Suspension, 32 mg, under this ANDA. You have notified the Agency that Natco Pharma Limited (Natco) 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 Natco for infringement of the '126 patent in the United States District Court for the District of New Jersey [Actelion Pharmaceuticals Ltd. and Actelion Pharmaceuticals US, Inc. v. Natco Pharma Limited, Civil Action No. 19-12984]. You have also notified the Agency that, on November 26, 2019, the court entered a Consent Judgment and Order with respect to Natco, which included that, “Natco agrees that Claims 4, 6, 7, 8, and 9 of the '126 patent are valid and enforceable with respect to the manufacture, use, offer for sale, sale, and importation of the Natco Product in the United States” and “Natco agrees that Natco's ANDA submission and the commercial manufacture, use, offer for sale, and/or sale within the United States, and/or importation into the United States of the Natco Product before the expiration of the '126 patent would constitute acts of infringement of Claims 4, 6, 7, 8, and 9 of the '126 patent”.¹ You have also provided a statement that Natco has obtained a license to the '126 patent; however, we note that 21 CFR 314.94(a)(12)(v) requires that, “[i]f the patent owner consents to approval of the ANDA (if otherwise eligible for approval) as of a specific date, the ANDA must contain a written statement from the patent owner that it has a licensing agreement with the applicant and that it consents to approval of the ANDA as of a specific date.” You have not submitted such a written statement from the patent owner. Therefore, because the Consent Judgment and Order is not a court order terminating the 30-month stay nor a court order of dismissal without a finding of infringement under 21 CFR 314.107(b)(3)(vii) or (viii), respectively, final approval cannot be granted until:

- the expiration of the 30-month period provided for in section 505(j)(5)(B)(iii) of the FD&C Act,
 - the date the court decides² that the '126 patent is invalid or not infringed (see sections 505(j)(5)(B)(iii)(I), (II), and (III) of the FD&C Act), or
 - the '126 patent has expired, and
- The Agency is assured there is no new information that would affect whether final approval should be granted.

Alternatively, you may also provide written consent to approval from the patent owner pursuant to 21 CFR 314.94(a)(12)(v) or from the patent owner or exclusive patent licensee pursuant to 21 CFR 314.107(b)(3)(vi).

In addition, the RLD upon which you have based your ANDA, Actelion's Tracleer Tablets for Oral Suspension, 32 mg, is subject to a period of exclusivity. As noted in the Orange Book, the ODE-161 exclusivity is scheduled to expire on September 5, 2024. Your ANDA contains a statement that you do not seek to market Bosentan Tablets for Oral Suspension, 32 mg, prior to the expiration of this exclusivity. Therefore, final approval also cannot be granted until the ODE-161 exclusivity has expired, currently September 5, 2024.

RISK EVALUATION AND MITIGATION STRATEGY REQUIREMENTS

Section 505-1 of the FD&C Act authorizes FDA to require the submission of a risk evaluation and mitigation strategy (REMS), if FDA determines that such a strategy is necessary to ensure that the benefits of the drug outweigh the risks [section 505-1(a)]. In accordance with section 505-1(i) of the FD&C Act, a drug that is the subject of an ANDA under section 505(j) is subject to certain elements of the REMS required for the applicable listed drug.

The details of the REMS requirements were outlined in our REMS notification letter dated April 4, 2019.

Your final proposed REMS, known as the Bosentan REMS Program, received on February 9, 2021, can be approved with your application.

The REMS consists of ETASU and an implementation system.

Prior to the submission of your amendment requesting final approval discussed below, you should contact Suzanne McLeod at Hikma, who has been identified as the current point of contact for the Bosentan REMS program, to obtain an updated version of the REMS and appended materials to submit to your application, **if applicable**. They can be reached at 614-241-4106 or smcleod@hikma.com.

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.

ANNUAL FACILITY FEES

The Generic Drug User Fee Amendments of 2012 (GDUFA) (Public Law 112-144, Title III) established certain provisions³ with respect to self-identification of facilities and payment of annual facility fees. Your ANDA identifies at least one facility that is subject to the self-identification requirement and payment of an annual facility fee. Self-identification must occur by June 1st of each year for the next fiscal year. Facility fees must be paid each year by the date specified in the *Federal Register* notice announcing facility fee amounts.

All finished dosage forms or active pharmaceutical ingredients manufactured in a facility that has not met its obligations to self-identify or to pay fees when they are due will be deemed misbranded. This means that it will be a violation of federal law to ship these products in interstate commerce or to import them into the United States. Such violations can result in prosecution of those responsible, injunctions, or seizures of

misbranded products. Products misbranded because of failure to self-identify or pay facility fees are subject to being denied entry into the United States.

In addition, we note that GDUFA requires that certain non-manufacturing sites and organizations listed in generic drug submissions comply with the self-identification requirement. The failure of any facility, site, or organization to comply with its obligation to self-identify and/or to pay fees when due may raise significant concerns about that site or organization and is a factor that may increase the likelihood of a site inspection prior to approval. FDA does not expect to give priority to completion of inspections that are required simply because facilities, sites, or organizations fail to comply with the law requiring self-identification or fee payment.

For further information on the status of this ANDA or upon submitting an amendment to the ANDA, please contact Olga Salis, Regulatory Project Manager, at (301) 796 - 0837.

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

¹ Consent Judgment and Order, *Actelion Pharmaceuticals Ltd. and Actelion Pharmaceuticals US, Inc. v. Natco Pharma Limited*, Civil Action No. 19-12984 (D.N.J., Nov. 26, 2019), at 2.

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.

³ Some of these provisions were amended by the Generic Drug User Fee Amendments of 2017 (GDUFA II) (Public Law 115-52, Title III).



Sarah
Kurtz

Digitally signed by Sarah Kurtz
Date: 4/01/2021 02:47:35PM
GUID: 54078879000a1b9e15dd31ed6f0343ca