

NDA 215644

COMPLETE RESPONSE

Sun Pharmaceutical Industries Limited C/O: Sun Pharmaceutical Industries, Inc. Attention: Juan Grijalva US Agent, Senior Manager 2 Independence Way Princeton, NJ 08540

Dear Juan Grijalva:

Please refer to your new drug applicatio	n (NDA)			(b) (4)	
	for	(b) (4)	(niloti	nib) capsules.	
We acknowledge receipt of your amend complete response to our		ted action le	(b) (4) tter.	, which constituted	а

We have completed our review of this application, and have determined that we cannot approve this application in its present form. We have described our reasons for this action below and, where possible, our recommendations to address these issues.

PRESCRIBING INFORMATION

(1) We reserve comment on the proposed labeling until the application is otherwise adequate. We encourage you to review the labeling review resources on the Prescription Drug Labeling Resources¹ and Pregnancy and Lactation Labeling Final Rule² websites, including regulations and related guidance documents and the Selected Requirements for Prescribing Information (SRPI) – a checklist of important format items from labeling regulations and guidances.

If you revise labeling, use the SRPI checklist to ensure that the Prescribing Information conforms with format items in regulations and guidances. Your response must include updated content of labeling [21 CFR 314.50(I)(1)] in structured product labeling (SPL) format as described at FDA.gov.³

https://www.fda.gov/drugs/laws-acts-and-rules/prescription-drug-labeling-resources

² https://www.fda.gov/drugs/labeling-information-drug-products/pregnancy-and-lactation-labeling-drugs-final-rule

³ http://www.fda.gov/ForIndustry/DataStandards/StructuredProductLabeling/default.htm

CARTON AND CONTAINER LABELING

(2) We reserve comment on the proposed labeling until the application is otherwise adequate.

MEDICATION GUIDE

(3) Add the following bolded statement or appropriate alternative to the carton and container labeling per 21 CFR 208.24(d): "ATTENTION PHARMACIST: Each patient is required to receive the enclosed Medication Guide."

PROPRIETARY NAME

(4) Please refer to correspondence dated, proposed proprietary name, the proposed proprietary name, the proposed proprietary name when you respond to the application deficiencies.

FACILITY INSPECTIONS

(5)	approval inspection of
	listed in this application, FDA
	conveyed deficiencies to the representative of the facility. The facility should
	provide satisfactory responses to these deficiencies to the FDA office indicated
	on the FDA 483 prior to your complete response to your application. Our
	determination that the facility's responses are satisfactory will depend on a
	finding that the facility has come into compliance with CGMP and has addressed
	any deficiencies specific to your application. You should coordinate with the
	facility for timely resolution of all inspection deficiencies, as well as to determine if
	any deficiencies may require updates to your application. Your complete
	response should include the date(s) of the facility's response(s) to the FDA Form
	483. Please refer to Compliance Program CP 7356.002 for guidance on post-
	inspection activities specific to CGMP compliance evaluation. FDA may
	determine that a CGMP reinspection and/or additional pre-approval inspection
	(PAI) is needed to confirm satisfactory resolution of inspection deficiencies
	before this application can be approved. If both CGMP and PAI reinspection are
	needed, the PAI coverage will generally occur following a determination that the
	facility is in compliance with CGMP.

SAFETY UPDATE

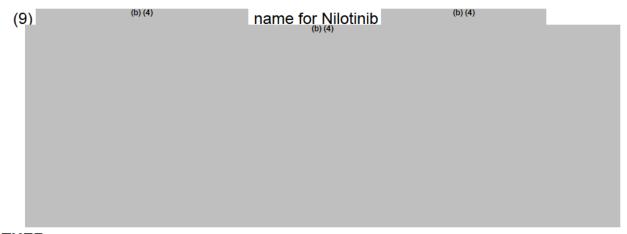
When you respond to the above deficiencies, include a safety update as described at 21 CFR 314.50(d)(5)(vi)(b). The safety update should include data from all nonclinical

U.S. Food and Drug Administration Silver Spring, MD 20993 www.fda.gov and clinical studies/trials of the drug under consideration regardless of indication, dosage form, or dose level.

- (1) Describe in detail any significant changes or findings in the safety profile.
- (2) When assembling the sections describing discontinuations due to adverse events, serious adverse events, and common adverse events, incorporate new safety data as follows:
 - Present new safety data from the studies/clinical trials for the proposed indication using the same format as in the original submission.
 - Present tabulations of the new safety data combined with the original application data.
 - Include tables that compare frequencies of adverse events in the original application with the retabulated frequencies described in the bullet above.
 - For indications other than the proposed indication, provide separate tables for the frequencies of adverse events occurring in clinical trials.
- (3) Present a retabulation of the reasons for premature trial discontinuation by incorporating the drop-outs from the newly completed trials. Describe any new trends or patterns identified.
- (4) Provide case report forms and narrative summaries for each subject who died during a clinical trial or who did not complete a trial because of an adverse event. In addition, provide narrative summaries for serious adverse events.
- (5) Describe any information that suggests a substantial change in the incidence of common, but less serious, adverse events between the new data and the original application data.
- (6) Provide updated exposure information for the clinical studies/trials (e.g., number of subjects, person time).
- (7) Provide a summary of worldwide experience on the safety of this drug. Include an updated estimate of use for drug marketed in other countries.
- (8) Provide English translations of current approved foreign labeling not previously submitted.

ADDITIONAL COMMENTS

We have the following comments/recommendations that are not approvability issues:



OTHER

Within one year after the date of this letter, you are required to resubmit or take other actions available under 21 CFR 314.110. If you do not take one of these actions, we may consider your lack of response a request to withdraw the application under 21 CFR 314.65. You may also request an extension of time in which to resubmit the application.

A resubmission must fully address all the deficiencies listed in this letter and should be clearly marked with "RESUBMISSION" in large font, bolded type at the beginning of the cover letter of the submission. The cover letter should clearly state that you consider this resubmission a complete response to the deficiencies outlined in this letter. A partial response to this letter will not be processed as a resubmission and will not start a new review cycle.

You may request a meeting or teleconference with us to discuss what steps you need to take before the application may be approved. If you wish to have such a meeting, submit your meeting request as described in the draft guidance for industry *Formal Meetings Between the FDA and Sponsors or Applicants of PDUFA Products*.

The drug product may not be legally marketed until you have been notified in writing that this application is approved.

If you have any questions, contact	(0) (4)
	Sincerely,
	{See appended electronic signature page}
	(b) (4)
	Center for Drug Evaluation and Research

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/

(b) (4)

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