



NDA 21-226/S-020

Abbott Laboratories
Attention: Raymond Votzmeyer
Associate Director, Global Pharmaceutical Regulatory Affairs
200 Abbott Park Road
Dept. RA76/Building AP30-1NE
Abbott Park, IL 60064-6157

Dear Mr. Votzmeyer:

Please refer to your supplemental new drug application dated December 28, 2006, received December 29, 2006, submitted under section 505(b) of the Federal Food, Drug, and Cosmetic Act for Kaletra[®] (133.3 mg lopinavir/33.3 mg ritonavir) Capsules.

We acknowledge receipt of your submissions dated May 18, 2007, June 22, 2007, June 27, 2007, and June 28, 2007.

This supplemental new drug application provides for revisions to the CLINICAL PHARMACOLOGY, Microbiology section, CLINICAL PHARMACOLOGY, Drug Interactions section, and the PRECAUTIONS section of the package insert.

We have completed our review of this application, as amended. This application is approved, effective on the date of this letter, for use as recommended in the agreed-upon labeling text.

Within 21 days of the date of this letter, submit content of labeling [21 CFR 314.50(l)] in structured product labeling (SPL) format, as described at <http://www.fda.gov/oc/datacouncil/spl.html>, that is identical in content to the enclosed labeling text. Upon receipt and verification, we will transmit that version to the National Library of Medicine for public dissemination.

The final printed labeling (FPL) must be identical to the enclosed labeling (text for the package insert and text for the patient package insert).

Please submit an electronic version of the FPL according to the guidance for industry document titled *Providing Regulatory Submissions in Electronic Format - NDA*. Alternatively, you may submit 20 paper copies of the FPL as soon as it is available but no more than 30 days after it is printed. Individually mount 15 of the copies on heavy-weight paper or similar material. For administrative purposes, designate this submission “**FPL for approved supplements NDA 21-226/S-020.**” Approval of this submission by FDA is not required before the labeling is used.

If you issue a letter communicating important information about this drug product (i.e., a “Dear Health Care Professional” letter), we request that you submit a copy of the letter to these NDAs and a copy to the following address:

MEDWATCH
Food and Drug Administration
5515 Security Lane
HFD-001, Suite 5100
Rockville, MD 20852

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 Karen Winestock, Regulatory Project Manager, at (301) 796-0834.

Sincerely,

{See appended electronic signature page}

Debra Birnkrant, M.D.
Director
Division of Antiviral Products
Office of Antimicrobial Products
Center for Drug Evaluation and Research

Enclosures: Package Insert
Patient Package Insert

**This is a representation of an electronic record that was signed electronically and
this page is the manifestation of the electronic signature.**

/s/

Jeffrey Murray

7/9/2007 08:02:42 AM