



NDA 20-716/S-008

Abbott Laboratories
200 Abbott Park Road
Dept RA76, BLDG AP30-1NE
Abbott Park, IL 60064-6157

Attention: David C. Ross, Pharm.D., MBA
Director, Regulatory Affairs

Dear Dr. Ross:

Please refer to your supplemental new drug application dated October 28, 2005, received October 30, 2005, submitted under section 505(b) of the Federal Food, Drug, and Cosmetic Act for Vicoprofen (hydrocodone bitartrate and ibuprofen) Tablets.

This supplemental new drug application was submitted in response to the Agency's letter dated June 14, 2005, requiring class labeling language for all non-selective non-steroidal anti-inflammatory drugs (NSAIDs), to include a boxed warning to address possible cardiovascular risks as well as known gastrointestinal risks, revised **CONTRAINDICATIONS, WARNINGS** and **PRECAUTIONS** sections of the package insert, and a **MedGuide** for NSAIDs.

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 agreed-upon labeling text.

The final printed labeling (FPL) must be identical, and include the revisions indicated, to the enclosed labeling text for the package insert and the MedGuide. The revisions were agreed upon in your October 13, 2006 email communication. These revisions are terms of the approval of this application.

Please submit an electronic version of the FPL according to the guidance for industry 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 supplement NDA 20-716/S-008.**" 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 this NDA 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 Sharon Turner-Rinehardt, Regulatory Project Manager, at (301) 827-2280.

Sincerely,

{See appended electronic signature page}

Bob Rappaport, M.D.
Director
Division of Anesthesia, Analgesia and
Rheumatology Products
Office of Drug Evaluation II
Center for Drug Evaluation and Research

Enclosure

**This is a representation of an electronic record that was signed electronically and
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/s/

Bob Rappaport
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