DEPARTMENT OF HEALTH & HUMAN SERVICES

Public Health Service

Food and Drug Administration Rockville, MD 20857

NDA 18-989/S-058

Wyeth Consumer Healthcare
Attention: Barbara Wolfe, PharmD.

Aggaginta Director Roman

Associate Director, Regulatory Affairs Five Giralda Farms Madison, New Jersey 07940

Dear Dr. Wolfe:

Please refer to your supplemental new drug application dated February 9, 2004, received February 12, 2004, submitted under section 505(b) of the Federal Food, Drug, and Cosmetic Act for Advil (200 mg ibuprofen) tablets.

This "Changes Being Effected" supplemental new drug application provides for the addition of new organ-specific warnings to the Drug Facts label.

We have completed our review of this supplemental new drug application. It is approved, effective on the date of this letter, for use as recommended in the final printed labeling (FPL), submitted on February 9, 2004, for the 10 count tablets (blister and bottle), 200 count tablets (carton and bottle), 100 count caplets (carton and bottle), and 225 count gel caplets (carton and bottle).

We are concerned about the need for organ-specific warnings for OTC drug products containing analgesic/antipyretic active ingredients. The proposed stomach bleeding warning is acceptable as interim language. However, please note that we will be providing guidance on wording and placement of organ-specific warnings in the labeling of drug products containing NSAIDs in the future.

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, HFD-410 FDA 5600 Fishers Lane Rockville, MD 20857

We remind you that you must comply with the requirements for an approved NDA set forth under 21 CFR 314.80 and 314.81.

If you have any questions, call Walter Ellenberg, Ph.D., Regulatory Project Manager, at (301) 827-2241.

Sincerely,

{See appended electronic signature page}

Charles Ganley, M.D.
Director
Division of Over-the-Counter Drug Products
Office of Drug Evaluation V
Center for Drug Evaluation and Research

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

/s/

Charles Ganley 7/16/04 03:22:04 PM