



DEPARTMENT OF HEALTH & HUMAN SERVICES

Public Health Service

ANDA 40-422

Food and Drug Administration
Rockville MD 20857

RECEIVED

FEB 11 2002

Barr Laboratories, Inc.
Attention: Christine Mundkur
2 Quaker Road
P.O. Box 2900
Pomona, NY 10970-0519

CHRISTINE MUNDKUR, ESQ.

FEB 11 2002

Dear Madam:

This is in reference to your abbreviated new drug application dated October 31, 2000, submitted pursuant to Section 505(j) of the Federal Food, Drug, and Cosmetic Act (Act), for Dextroamphetamine Saccharate, Amphetamine Aspartate, Dextroamphetamine Sulfate, and Amphetamine Sulfate Tablets, 5 mg, 10 mg, 20 mg, and 30 mg. Each drug product strength represents the total of equal amounts of the four component salts.

Reference is also made to your amendments dated May 17, June 8, July 11, August 31, September 4, September 21, October 5, and November 14, 2001.

We have completed the review of this abbreviated application and have concluded that the drug is safe and effective for use as recommended in the submitted labeling. Accordingly the application is approved. The Division of Bioequivalence has determined your tablets containing combined salts of Dextroamphetamine Saccharate, Amphetamine Aspartate, Dextroamphetamine Sulfate, and Amphetamine Sulfate Tablets, 5 mg, 10 mg, 20 mg, and 30 mg, to be bioequivalent and, therefore, therapeutically equivalent to the listed drug (Adderal[®] Tablets, 5 mg, 10 mg, 20 mg, and 30 mg, respectively, of Shire Laboratories Inc.). Your dissolution testing should be incorporated into the stability and quality control program using the same method proposed in your application.

Under Section 506A of the Act, certain changes in the conditions described in this abbreviated application require an approved supplemental application before the change may be made.

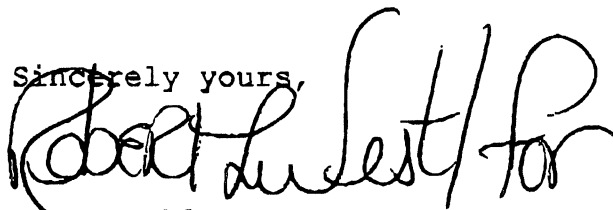
Post-marketing reporting requirements for this abbreviated application are set forth in 21 CFR 314.80-81 and 314.98. The Office of Generic Drugs should be advised of any change in the marketing status of this drug.

We request that you submit, in duplicate, any proposed advertising or promotional copy that you intend to use in your initial advertising or promotional campaigns. Please submit all proposed materials in draft or mock-up form, not final print. Submit both copies together with a copy of the proposed or final printed labeling to the Division of Drug Marketing, Advertising, and Communications (HFD-40). Please do not use Form FD-2253 (Transmittal of Advertisements and Promotional Labeling for Drugs for Human Use) for this initial submission.

We call your attention to 21 CFR 314.81(b)(3) which requires that materials for any subsequent advertising or promotional campaign be submitted to our Division of Drug Marketing, Advertising, and Communications (HFD-40) with a completed Form FD-2253 at the time of their initial use.

Validation of the regulatory methods has not been completed. It is the policy of the Office not to withhold approval until the validation is complete. We acknowledge your commitment to satisfactorily resolve any deficiencies that may be identified.

Sincerely yours,



Gary Buehler

Director

Office of Generic Drugs

Center for Drug Evaluation and Research