

**CENTER FOR DRUG EVALUATION AND
RESEARCH**

APPLICATION NUMBER:

40372

APPROVAL LETTER

ANDA 40-372

JUN 8 2000

Abbott Laboratories
Attention: Jessie Y. Lee
200 Abbott Park Road
Dept. 389, Bldg. AP30
Abbott Park, IL 60064-6157

Dear Madam:

This is in reference to your abbreviated new drug application dated May 28, 1999, submitted pursuant to Section 505(j) of the Federal Food, Drug, and Cosmetic Act (Act), for Promethazine Hydrochloride Injection USP, 25 mg/mL and 50 mg/mL (Carpject).

Reference is also made to your amendments dated January 18, April 14, May 19, and May 22, 2000.

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 Promethazine Hydrochloride Injection USP, 25 mg/mL and 50 mg/mL (Carpject) to be bioequivalent and, therefore, therapeutically equivalent to the listed drug (Phenergan Injection, 25 mg/mL and 50 mg/mL of Wyeth Ayerst Laboratories, Inc.).

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 which 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

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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.

Sincerely yours,

[S]
Gary Buehler 6/8/00
Acting Director
Office of Generic Drugs
Center for Drug Evaluation and Research

cc: ANDA 40-372
Division File
FIELD COPY
HFD-610/R.West
HFD-92
HFD-210/B.Poole
HFD-330/
HFD-205/

W. [Signature] 6/7/00

APPROVAL