

**CENTER FOR DRUG EVALUATION AND  
RESEARCH**

*APPLICATION NUMBER:*

**75-547**

**APPROVAL LETTER**

APR 2 2001

Bedford Laboratories  
A Division of BenVenue Laboratories, Inc.  
Attention: Shahid Ahmed  
300 Northfield Road  
Bedford, Ohio 44146

Dear Sir:

This is in reference to your abbreviated new drug application dated December 29, 1998, submitted pursuant to Section 505(j) of the Federal Food, Drug, and Cosmetic Act (Act), for Thiotepa for Injection, 15 mg/vial.

Reference is also made to your amendments dated February 2, February 26 and March 1, 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 Thiotepa for Injection, 15 mg/vial, to be bioequivalent and, therefore, therapeutically equivalent to the listed drug (Thioplex<sup>®</sup> Injection, 15 mg/vial of Immunex Corp).

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 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,



Gary Buehler 4/2/01  
Acting Director  
Office of Generic Drugs  
Center for Drug Evaluation and Research

ANDA 75-547

Bedford Laboratories  
Attention: Shahid Ahmed  
270 Northfield Road  
Bedford, OH 44146  
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FEB 9 1999

Dear Sir:

We acknowledge the receipt of your abbreviated new drug application submitted pursuant to Section 505(j) of the Federal Food, Drug and Cosmetic Act.

Reference is made to the telephone conversation dated January 28, 1999 and your correspondence dated February 4, 1999.

NAME OF DRUG: Thiotepa for Injection, 15 mg/vial

DATE OF APPLICATION: December 29, 1998

DATE (RECEIVED) ACCEPTABLE FOR FILING: December 31, 1998

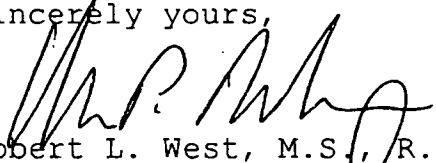
We will correspond with you further after we have had the opportunity to review the application.

Please identify any communications concerning this application with the ANDA number shown above.

Should you have questions concerning this application, contact:

Denise Huie  
Project Manager  
(301) 827-5848

Sincerely yours,

  
Robert L. West, M.S., R.Ph.  
Director  
Division of Labeling and Program Support  
Office of Generic Drugs  
Center for Drug Evaluation and Research