

ANDA 65-062

November 30, 2000

Ranbaxy Pharmaceuticals Inc.  
Attention: Shirley Ternyik, U.S. Agent for  
Ranbaxy Laboratories Limited  
600 College Road East  
Princeton, NJ 08540

Dear Madam:

This is in reference to your abbreviated new drug application dated December 27, 1999, submitted pursuant to Section 505(j) of the Federal Food, Drug, and Cosmetic Act (Act), for Minocycline Hydrochloride Capsules USP, 50 mg(base), 75 mg(base), and 100 mg (base). We note that this product is subject to the exception provisions of Section 125(d)(2) of Title I of the Food and Drug Administration Modernization Act of 1997.

Reference is also made to your amendments dated July 27, October 13, November 8, and November 16, 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 Minocycline Hydrochloride Capsules USP, 50 mg (base), 75 mg(base), and 100 mg(base), to be bioequivalent and, therefore, therapeutically equivalent to the listed drug (Minocycline Hydrochloride Capsules USP, 50 mg(base), 75 mg(base), and 100 mg(base), respectively, of Danbury Pharmacal, 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 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  
Acting Director  
Office of Generic Drugs  
Center for Drug Evaluation and Research