

**CENTER FOR DRUG EVALUATION AND
RESEARCH**

APPLICATION NUMBER:

75-271

APPROVAL LETTER

JAN 18 2000

Zenith Goldline Pharmaceuticals, Inc.
Attention: Jason Gross, Pharm. D.
U.S. Agent for: Steripak Limited
140 Legrand Avenue
Northvale, NJ 07647-2485

Dear Sir:

This is in reference to your abbreviated new drug application dated December 11, 1997, submitted pursuant to Section 505(j) of the Federal Food, Drug, and Cosmetic Act (Act), for Cromolyn Sodium Inhalation Solution USP, 10 mg/mL, packaged in 2 mL unit-dose vials.

Reference is also made to your amendments dated June 24, July 29, November 15, December 1, and December 17, 1999.

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 Cromolyn Sodium Inhalation Solution USP, 10 mg/mL, to be bioequivalent and, therefore, therapeutically equivalent to the listed drug (Intal[®] Nebulizer Solution, 10mg/mL, of Rhone Poulenc Rorer Pharmaceuticals 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 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,


Douglas L. Sporn 1/18/00
Director

Office of Generic Drugs
Center for Drug Evaluation and Research