DEPARTMENT OF HEALTH & HUMAN SERVICES

Public Health Service

Food and Drug Administration
Rockville MD 20857

ANDA 74-185

MAY 31 1995

Lemmon Company
Attention: Deborah Jaskot
650 Cathill Road
Sellersville, PA 18960

Dear Ms. Jaskot:

This is in reference to your abbreviated new drug application dated March 4, 1992, submitted pursuant to Section 505(j) of the Federal Food, Drug, and Cosmetic Act, for Diltiazem Hydrochloride Tablets USP, 30 mg, 60 mg, 90 mg and 120 mg.

Reference is also made to your amendments dated May 5, 1995 and May 30, 1995.

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 Diltiazem Hydrochloride Tablets, 30 mg, 60 mg, 90 mg and 120 mg, to be bioequivalent and, therefore, therapeutically equivalent to those of the listed drug (CARDIZEM® Tablets, 30 mg, 60 mg, 90 mg, and 120 mg, respectively, of Marion Merrell Dow Inc.). Your dissolution testing should be incorporated into the stability and quality control program using the same method proposed in your application.

Under 21 CFR 314.70, 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. 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-240). 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-240) with a completed Form FD-2253 at the time of their initial use.

Sincerely yours,

D. L. Sporn

Douglas L. Sporn
Acting Director
Office of Generic Drugs
Center for Drug Evaluation and Research