ANDA 74-555

SEP 30, 1998

Copley Pharmaceutical, Inc.
Attention: Isidoro Nudelman
25 John Road
Canton, MA 02021

Dear Sir:

This is in reference to your abbreviated new drug application dated October 3, 1994, submitted pursuant to Section 505(j) of the Federal Food, Drug, and Cosmetic Act, for Cholestyramine for Oral Suspension, USP (Light), 4 g Resin/Packet and 4 g Resin/Scoopful.

Reference is also made to your amendments dated Reference is also made to your amendments dated August 23, 1996; November 19, 1997; February 26, July 1, 6, and 28, 1998.

This corrects our approval letter of September 30, 1998.

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 Cholestyramine for Oral Suspension, USP (Light), 4 g Resin/Packet and 4 g Resin/Scoopful to be bioequivalent and, therefore, therapeutically equivalent to the listed drug (Questran Powder, 4 g resin per packet and 4 g resin per scoopful,of Bristol Myers Co.). The binding capacity assay described in USP 23 should be incorporated into your stability and quality control program.

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 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
Director
Office of Generic Drugs
Center for Drug Evaluation and Research