



NDA 20-369/S-009 & S-010

Alcon Laboratories, Inc.
c/o Alcon Research, Ltd.
Attention: Richard O. Reese
Regulatory Affairs
6201 South Freeway
Fort Worth, TX 76134

Dear Mr. Reese:

Please refer to your supplemental new drug applications dated February 6, 2004, received February 9, 2004, submitted under 505(b) the Federal Food, Drug, and Cosmetic Act for Ciloxan (ciprofloxacin hydrochloride ophthalmic ointment) 0.3%.

These "Changes Being Effected" supplemental new drug applications provide for a change to the tamper evident feature to the drug product outer packaging and labeling changes.

We completed our review of these supplemental new drug applications. They are approved, effective on the date of this letter, for use as recommended in the final printed labeling (FPL), including the package insert, carton and container labels, received on February 9, 2004.

However, if a future labeling supplement is submitted, please revise the product carton to include a precautionary statement as in the package insert, "Do not use the product if the imprinted carton seals have been damaged or removed."

If you issue a letter communicating important information about this drug product (i.e., a "Dear Health Care Professional" letter), we request that you submit a copy of the letter to this NDA and a copy to the following address:

MEDWATCH, HFD-410
FDA
5600 Fishers Lane
Rockville, MD 20857

We remind you that you must comply with the requirements for an approved NDA set forth under 21 CFR 314.80 and 314.81.

If you have any questions, call Lori M. Gorski, Project Manager, at (301) 827-2090.

Sincerely,

{See appended electronic signature page}

Wiley A. Chambers, M.D.
Deputy Director
Division of Anti-Inflammatory, Analgesic,
and Ophthalmic Drug Products, HFD-550
Office of Drug Evaluation V
Center for Drug Evaluation and Research

Enclosure

**This is a representation of an electronic record that was signed electronically and
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/s/

Wiley Chambers

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