



NDA 50-555/S-017, S-018 & S-019

Alcon Laboratories, Inc.  
c/o Alcon Research, Ltd.  
Attention: Sarah J. Cantrell  
Manager, Regulatory Affairs  
6201 South Freeway  
Fort Worth, Texas 76134-2099

Dear Ms. Cantrell:

Please refer to your supplemental new drug applications submitted under the Federal Food, Drug, and Cosmetic Act for Tobrex (tobramycin ophthalmic ointment) 0.3%.

Supplement	Originally Dated	Approvable Letter Dated	Amendments Dated
S-017	July 13, 2001	February 22, 2002	March 7, 2002 December 10, 2002
S-018	June 4, 2002	October 18, 2002	November 14, 2002
S-019	June 4, 2002	October 18, 2002	November 14, 2002

These supplemental new drug applications provide for the addition of the Puurs, Belgium facility as an alternate manufacturing site for the drug product; (b)(4)-----as an alternate drug substance (b)(4)----- site; revised tobramycin and drug product specifications; an alternate container closure; and revised labeling.

We completed our review of these supplemental new drug applications, as amended. They are approved, effective on the date of this letter, for use as recommended in the attached package insert submitted as final printed labeling (FPL) on November 14, 2002.

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, HF-2  
FDA  
5600 Fishers Lane  
Rockville, MD 20857

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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}*

Linda L. Ng, Ph.D.  
Chemistry Team Leader, for the  
Division of Anti-Inflammatory, Analgesic  
and Ophthalmic Drug Products, HFD-550  
DNDC III, Office of New Drug Chemistry  
Center for Drug Evaluation and Research

Enclosure

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/s/

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Linda Ng  
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