



NDA 50-679/S-023

Bristol-Myers Squibb Company  
Attention: David L. Silberstein  
Associate Director, Life Cycle Management  
P.O. Box 4000  
Princeton, NJ 08543-4000

Dear Mr. Silberstein:

Please refer to your supplemental new drug application dated January 14, 2004, received January 16, 2004, submitted under section 505(b) of the Federal Food, Drug, and Cosmetic Act for Maxipime<sup>®</sup> (cefepime hydrochloride) for Injection. This application is subject to the exemption provisions contained in section 125(d)(2) of Title I of the FDA Modernization Act of 1997.

This supplemental application, submitted as "Supplement - Changes Being Effected in 0 days," proposes the following change to be in compliance with the "Labeling Requirements for Systemic Antibacterial Drug Products Intended for Human Use" Final Rule, as published in the Federal Register, Vol. 68, No. 25, February 6, 2003.

We have completed our review of this application. It is approved, effective on the date of this letter, for use as recommended in the final printed labeling (FPL) submitted on January 16, 2004.

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

We remind you that you must comply with reporting requirements for an approved NDA (21 CFR 314.80 and 314.81).

If you have any questions, call LT Raquel Peat, Regulatory Health Project Manager, at (301) 827-2125.

Sincerely,

*{See appended electronic signature page}*

Janice M. Soreth, M.D.

Director

Division of Anti-Infective Drug Products

Office of Drug Evaluation IV

Center for Drug Evaluation and Research

Enclosure: Labeling

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/s/

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Janice Soreth

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