



NDA 50-671/S-006

Baxter Healthcare Corporation
Attention: Marcia Marconi
Vice President, Regulatory Affairs
Route 120 and Wilson Road; RLT-10
Round Lake, IL 60073

Dear Ms. Marconi:

Please refer to your supplemental new drug application dated September 22, 2003, received September 23, 2003, submitted under section 505(b) of the Federal Food, Drug, and Cosmetic Act for Vancocin® HCL (vancomycin injection, USP) in Galaxy® Plastic Container, PL 2040. We note that this application is subject to the exemption provisions contained in section 125(d)(2) of Title I of the FDA Modernization Act of 1997.

We acknowledge receipt of your submission dated October 10, 2003.

This supplemental application, submitted as a "Supplement - Changes Being Effected in 0 days" supplement, proposes the following change to be in compliance with the systemic antibacterial drug products labeling regulations as found in 21 CFR 201.24 and the addition of contraindication for corn allergy.

We have completed our review of this application, as amended. It is approved, effective on the date of this letter, for use as recommended in the final printed labeling (FPL) submitted on September 22, 2003.

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 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

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

Janice Soreth

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