Dear Mr. Clark:

Please refer to your supplemental new drug applications dated December 9, 2003, received December 10, 2003, submitted under section 505(b) of the Federal Food, Drug, and Cosmetic Act for Vantin® (cefpodoxime proxetil) Tablets, NDA 50-674/S-013 and Vantin® (cefpodoxime proxetil) Oral Suspension, NDA 50-675/S-016. We note that these applications are subject to the exemption provisions contained in section 125(d)(2) of Title I of the FDA Modernization Act of 1997.

These supplemental applications, 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 these supplemental new drug applications, as amended. They are approved, effective on the date of this letter, for use as recommended in the final printed labeling (FPL) submitted on December 10, 2003.

If a letter communicating important information about this drug product (i.e., a "Dear Health Care Professional" letter) is issued to physicians and others responsible for patient care, 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 the requirements for an approved NDA set forth under 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 this page is the manifestation of the electronic signature.

/s/

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Janice Soreth
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