



NDA 21-892/S-001

Salix Pharmaceuticals, Inc.
Attention: Jill Kompa, M.S., RAC
Director, Regulatory Affairs
1700 Perimeter Park Drive
Morrisville, NC 27560

Dear Ms. Kompa:

Please refer to your supplemental new drug application dated March 19, 2007, received March 21, 2007, submitted under section 505(b) of the Federal Food, Drug, and Cosmetic Act for OsmoPrep (sodium phosphate monobasic monohydrate, USP & sodium phosphate dibasic anhydrous, USP) Tablets.

We also acknowledge receipt of your submission dated March 23, 2007.

This "Changes Being Effected" supplemental new drug application provides for revisions to the Package Insert (PI), including the addition of "Postmarketing Experience" subsection to the "ADVERSE REACTIONS" section as well as some editorial changes.

We completed our review of this application, as amended. This application is approved, effective on the date of this letter, for use as recommended in the agreed-upon labeling text.

The final printed labeling (FPL) must be identical to the package insert submitted on March 19, 2007.

Please submit an electronic version of the FPL according to the guidance for industry titled *Providing Regulatory Submissions in Electronic Format - NDA*. Alternatively, you may submit 20 paper copies of the FPL as soon as it is available but no more than 30 days after it is printed. Individually mount 15 of the copies on heavy-weight paper or similar material. For administrative purposes, designate this submission "**FPL for approved supplement NDA 21-892/S-001.**" Approval of this submission by FDA is not required before the labeling is used.

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
Food and Drug Administration
5515 Security Lane
HFD-001, Suite 5100
Rockville, MD 20852

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 Matthew Scherer, Regulatory Project Manager, at (301) 796-2307.

Sincerely,

{See appended electronic signature page}

Joyce Korvick, M.D., M.P.H.
Deputy Director
Division of Gastroenterology Products
Office of Drug Evaluation III
Center for Drug Evaluation and Research

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

Joyce Korvick
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