



NDA 21-520 S-016

Eli Lilly and Company
Attention: Christine A. Phillips, Ph.D., RAC
Manager, U.S. Regulatory Affairs
Lilly Corporate Center
Indianapolis, IN 46285

Dear Dr. Phillips:

Please refer to your supplemental new drug applications dated and received January 10, 2008, submitted under section 505(b) of the Federal Food, Drug, and Cosmetic Act for Symbyax (olanzapine and fluoxetine HCl) Capsules.

This "Changes Being Effected" supplemental new drug application provides for new language to inform clinicians of the possible risk factors for and signs and symptoms of hyponatremia and to standardize the language across the SSRIs and SNRIs, as requested in the Agency letter of August 7, 2007. Specifically:

1. Revisions under the PRECAUTIONS section, Hyponatremia subsection.
2. Revisions under the Geriatric Use section.

We have completed our review of these supplemental new drug applications and they are approved effective on the date of this letter, for use as recommended in the final printed labeling (FPL) submitted on January 10, 2008 (copy attached).

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 the requirements for an approved NDA set forth under 21 CFR 314.80 and 314.81.

NDA 21-520 S-016

Page 2

If you have any questions, email CAPT Steven D. Hardeman, R.Ph., Chief, Project Management Staff, at Steven.Hardeman@FDA.HHS.GOV.

Sincerely,

{See appended electronic signature page}

Thomas Laughren, M.D.
Director
Division of Psychiatry Products
Office of Drug Evaluation I
Center for Drug Evaluation and Research

**This is a representation of an electronic record that was signed electronically and
this page is the manifestation of the electronic signature.**

/s/

Thomas Laughren
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