



NDA 21411/S-047

**SUPPLEMENT APPROVAL**

Eli Lilly and Company  
Attention: Marija Popovic, Ph.D.  
Consultant, US Regulatory Affairs  
Lilly Corporate Center, Drop Code 2543  
Indianapolis, IN 46285

Dear Dr. Popovic:

Please refer to your Supplemental New Drug Application (sNDA) dated April 02, 2015, submitted under section 505(b) of the Federal Food, Drug, and Cosmetic Act (FDCA) for Strattera (atomoxetine hydrochloride) 10 mg, 18 mg, 25 mg, 40 mg, 60 mg, 80 mg, and 100 mg capsules.

We acknowledge receipt of your amendment dated April 03, 2015.

We also refer to our letter dated February 2, 2015 notifying you, under Section 505(o)(4) of the FDCA, of new safety information that we believe should be included in the labeling for Strattera (atomoxetine hydrochloride). This information pertains to the association between the use of stimulants used to treat Attention Deficit Hyperactivity Disorder (ADHD) and rhabdomyolysis.

This supplemental new drug application provides for the addition of a new subsection "Musculoskeletal" to the Postmarketing Spontaneous Reports section of labeling, with "Rhabdomyolysis" added as a new adverse reaction term in this subsection consistent with our February 2, 2015 letter.

**APPROVAL & LABELING**

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

**CONTENT OF LABELING**

As soon as possible, but no later than 14 days from the date of this letter, submit the content of labeling [21 CFR 314.50(1)] in structured product labeling (SPL) format using the FDA automated drug registration and listing system (eLIST), as described at <http://www.fda.gov/ForIndustry/DataStandards/StructuredProductLabeling/default.htm>. Content of labeling must be identical to the enclosed labeling (text for the package insert, Medication

Guide), with the addition of any labeling changes in pending “Changes Being Effected” (CBE) supplements, as well as annual reportable changes not included in the enclosed labeling.

Information on submitting SPL files using eList may be found in the guidance for industry titled “SPL Standard for Content of Labeling Technical Qs and As at <http://www.fda.gov/downloads/DrugsGuidanceComplianceRegulatoryInformation/Guidances/UCM072392.pdf>

The SPL will be accessible from publicly available labeling repositories.

Also within 14 days, amend all pending supplemental applications that includes labeling changes for this NDA, including CBE supplements for which FDA has not yet issued an action letter, with the content of labeling [21 CFR 314.50(l)(1)(i)] in MS Word format, that includes the changes approved in this supplemental application, as well as annual reportable changes and annotate each change. To facilitate review of your submission, provide a highlighted or marked-up copy that shows all changes, as well as a clean Microsoft Word version. The marked-up copy should provide appropriate annotations, including supplement number(s) and annual report date(s).

## **REPORTING REQUIREMENTS**

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, contact Hiren Patel, Regulatory Project Manager, at [hiren.patel@fda.hhs.gov](mailto:hiren.patel@fda.hhs.gov).

Sincerely,

*{See appended electronic signature page}*

Victor Crentsil, MD, MHS  
Deputy Director for Safety  
Division of Psychiatry Products  
Office of Drug Evaluation I  
Center for Drug Evaluation and Research

ENCLOSURE:  
Content of Labeling

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**This is a representation of an electronic record that was signed electronically and this page is the manifestation of the electronic signature.**  
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/s/  
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VICTOR CRENTSIL  
04/17/2015