



NDA 021540/S-034

SUPPLEMENT APPROVAL

Pfizer Ireland Pharmaceuticals
C/O Pfizer Inc
Attention: Marcio de Godoy, MBA, PharmD, PhD
Global Regulatory Lead, Worldwide Safety and Regulation
235 East 42nd Street
New York, NY 10017

Dear Dr. Godoy:

Please refer to your Supplemental New Drug Application (sNDA) dated November 21, 2013, received November 21, 2013, submitted under section 505(b)(1) of the Federal Food, Drug, and Cosmetic Act (FDCA) for Caduet (atorvastatin/amlodipine) 5/10, 10/10, 5/20, 10/20, 5/40, 10/40, 5/80 and 10/80 mg.

This "Changes Being Effected" supplemental new drug application provides for corrections to the Section **12.3 Pharmacokinetics** and Section **14.11 Atorvastatin for Heterozygous Familial Hypercholesterolemia in Pediatric Patients**. Additional minor editorial changes were also made.

The following content changes were made;

1. Update table 5 in section **12.3 Pharmacokinetics** to correct the information for cimetidine, as this should be Cimetidine 300 mg QID, 2 weeks (changed from Cimetidine 300 mg QD, 4 weeks)
2. Update section **14.11 Atorvastatin for Heterozygous Familial Hypercholesterolemia in Pediatric Patients** with the number of atorvastatin-treated patients who required up-titration to 20 mg after Week 4 during the double-blind phase from 80 (57.1%) to 78 (55.7%)

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(l)] 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), 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, please call Michael Monteleone, Regulatory Project Manager, at (301) 796-1952.

Sincerely,

{See appended electronic signature page}

Mary Ross Southworth, PharmD
Deputy Director for Safety
Division of Cardiovascular and Renal Products
Office of Drug Evaluation I
Center for Drug Evaluation and Research

ENCLOSURE(S):

Content of Labeling

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

/s/

MARY R SOUTHWORTH
05/22/2014