



NDA 004782/S-162
NDA 004782/S-164
NDA 004782/S-167

SUPPLEMENT APPROVAL

Wyeth Pharmaceuticals, Inc.
c/o Pfizer Inc.
Attention: Ursula Campbell
Senior Director, WRS
235 East 42nd Street
New York, NY 10017

Dear Ms. Campbell:

Please refer to your Supplemental New Drug Applications (sNDAs) dated and received December 3, 2009, for S-162; March 2, 2010, for S-164; and June 23, 2010, for S-167 submitted under section 505(b) of the Federal Food, Drug, and Cosmetic Act (FDCA) for PREMARIN® (conjugated estrogens tablets, USP).

We acknowledge receipt of your amendments dated December 15, 2010, February 18, 2011, May 18, 2011, and June 28, 2011.

These "Prior Approval" (S-162) and "Changes Being Effected" (S-164 and S-167) supplemental new drug applications provide for revisions to the PATIENT INFORMATION section of the Patient Package Insert, a change to the WARNINGS section of the labeling, and changes to the CONTRAINDICATIONS and PRECAUTIONS sections of the labeling, respectively. These revisions provide information about the appearance of tablet in stool, an addition of a new WARNINGS of angioedema, an addition of a new CONTRAINDICATIONS of known thrombophilic disorders and a modified statement in the PRECAUTIONS section. Selected sections of the Prescribing Information and Patient Package Insert were revised to reflect current recommended estrogen-class labeling.

We have completed our review of these supplemental applications, as amended. They are approved, effective on the date of this letter, for use as recommended in the enclosed, agreed-upon labeling text.

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, text for the patient 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 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).

REQUIRED PEDIATRIC ASSESSMENTS

Under the Pediatric Research Equity Act (PREA) (21 U.S.C. 355c), all applications for new active ingredients, new indications, new dosage forms, new dosing regimens, or new routes of administration are required to contain an assessment of the safety and effectiveness of the product for the claimed indication(s) in pediatric patients unless this requirement is waived, deferred, or inapplicable.

Because none of these criteria apply to your application, you are exempt from this requirement.

PROMOTIONAL MATERIALS

All promotional materials that include representations about your drug product must be promptly revised to be consistent with the labeling changes approved in this supplement, including any new safety information [21 CFR 314.70(a)(4)]. The revisions in your promotional materials should include prominent disclosure of the important new safety information that appears in the revised package labeling. Within 7 days of receipt of this letter, submit your statement of intent to comply with 21 CFR 314.70(a)(4) to the address above or by fax to 301-847-8444.

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, call Samantha Bell, Regulatory Project Manager, at (301) 796-9687.

Sincerely,

{See appended electronic signature page}

Christine P. Nguyen, M.D.
Acting Deputy Director of Safety
Division of Reproductive and Urologic Products
Office of Drug Evaluation III
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/

CHRISTINE P NGUYEN
10/28/2011