



NDA 022272/S-017

SUPPLEMENT APPROVAL

Purdue Pharma, L.P.
One Stamford Forum
Stamford, CT 06901-3431

Attention: Beth Connelly
Associate Director, Regulatory Affairs

Dear Ms. Connelly:

Please refer to your Supplemental New Drug Application (sNDA) dated October 16, 2012, received October 19, 2012, submitted under section 505(b) of the Federal Food, Drug, and Cosmetic Act (FDCA) for OXYCONTIN (oxycodone hydrochloride controlled-release) Tablets.

We acknowledge receipt of your amendments dated December 13, 2012, and March 29, 2013.

This supplemental new drug application proposes modifications to the approved risk evaluation and mitigation strategy (REMS) for OXYCONTIN.

We have completed our review of this supplemental application, as amended. It is approved, effective on the date of this letter.

RISK EVALUATION AND MITIGATION STRATEGY REQUIREMENTS

The REMS for OXYCONTIN was originally approved on April 5, 2010, and modified on July 9, 2012. The REMS consists of a Medication Guide, elements to assure safe use, and a timetable for submission of assessments of the REMS. Your proposed modifications to the REMS consist of:

- Revisions to *Section VI. Specific Drug information for ER/LA Opioid Analgesic Products* of the FDA Blueprint
- Revisions to the REMS Website, including the landing page and the webpage listing covered products under the REMS program
- Revisions to individual product Medication Guides for relevant drugs
- Revision to the REMS document to remove ANDA holders from the Timetable for Submission of Assessments

Your proposed modified REMS, submitted on March 29, 2013, and appended to this letter, is approved.

The timetable for submission of assessments of the REMS will remain the same as that approved on July 9, 2012. There are no changes to the REMS assessment plan described in our July 9, 2012 letter.

This REMS uses a single, shared system for the elements to assure safe use and the REMS assessments. This single, shared system, known as the ER/LA Opioid Analgesic REMS Program, currently includes the products listed in Appendix 1. Other products may be added in the future if additional NDAs or ANDAs are approved.

We remind you that section 505-1(f)(8) of FDCA prohibits holders of an approved covered application with elements to assure safe use from using any element to block or delay approval of an application under section 505(b)(2) or (j). A violation of this provision in 505-1(f) could result in enforcement action.

The requirements for assessments of an approved REMS under section 505-1(g)(3) include with respect to each goal included in the strategy, an assessment of the extent to which the approved strategy, including each element of the strategy, is meeting the goal or whether one or more such goals or such elements should be modified.

In addition to the assessments submitted according to the timetable included in the approved REMS, you may propose a modification to the approved REMS when you submit a supplemental application for a new indication for use as described in section 505-1(g)(2)(A) of the FDCA.

If the assessment instruments and methodology for your REMS assessments are not included in the REMS supporting document, or if you propose changes to the submitted assessment instruments or methodology, you should update the REMS supporting document to include specific assessment instrument and methodology information at least 90 days before the assessments will be conducted. Updates to the REMS supporting document may be included in a new document that references previous REMS supporting document submission(s) for unchanged portions. Alternatively, updates may be made by modifying the complete previous REMS supporting document, with all changes marked and highlighted. Prominently identify the submission containing the assessment instruments and methodology with the following wording in bold capital letters at the top of the first page of the submission:

**NDA 022272 REMS CORRESPONDENCE
(insert concise description of content in bold capital letters, e.g.,
UPDATE TO REMS SUPPORTING DOCUMENT - ASSESSMENT
METHODOLOGY)**

An authorized generic drug under this NDA must have an approved REMS prior to marketing. Should you decide to market, sell, or distribute an authorized generic drug under this NDA, contact us to discuss what will be required in the authorized generic drug REMS submission.

Prominently identify the submission containing the REMS assessments or proposed modifications of the REMS with the following wording in bold capital letters at the top of the first page of the submission as appropriate:

NDA 022272 REMS ASSESSMENT

**NEW SUPPLEMENT FOR NDA 022272
PROPOSED REMS MODIFICATION**

**NEW SUPPLEMENT (NEW INDICATION FOR USE)
FOR NDA 022272
REMS ASSESSMENT
PROPOSED REMS MODIFICATION (if included)**

If you do not submit electronically, please send 5 copies of REMS-related submissions.

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 Lisa E. Basham, Senior Regulatory Health Project Manager, at (301) 796-1175.

Sincerely,

{See appended electronic signature page}

Judith A. Racoosin, M.D., M.P.H.
Deputy Director for Safety
Division of Anesthesia, Analgesia,
and Addiction Products
Office of Drug Evaluation II
Center for Drug Evaluation and Research

ENCLOSURE:
REMS

Appendix 1 List of applications

NDA 021260 AVINZA (morphine sulfate) extended-release capsules and its generic equivalent

NDA 021306 BUTRANS (buprenorphine) Transdermal System for transdermal administration

NDA 006134 DOLOPHINE (methadone hydrochloride) tablets and its generic equivalents

ANDA 087997 Methadone Oral Solution and its generic equivalents ANDA 087393
Methadone Oral Solution and its generic equivalents ANDA 089897
Methadone Oral Concentrate

NDA 019813 DURAGESIC (Fentanyl Transdermal System) for transdermal administration and its generic equivalents

NDA 022321 EMBEDA (morphine sulfate and naltrexone hydrochloride) extended- release capsules

NDA 021217 EXALGO (hydromorphone HCl) Extended-Release Tablets

NDA 020616 KADIAN (morphine sulfate) extended-release capsules and its generic equivalent

NDA 019516 MS CONTIN (morphine sulfate) controlled-release tablets and its generic equivalents

NDA 200533 NUCYNTA ER (tapentadol) extended-release oral tablets

NDA 201655 OPANA ER (oxymorphone hydrochloride) Extended-Release tablets

NDA 021610 OPANA ER (oxymorphone hydrochloride) Extended-Release tablets and its generic equivalents

NDA 022272 OXYCONTIN (oxycodone hydrochloride controlled-release) Tablets

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/s/

JUDITH A RACOOSIN
04/15/2013