
**UNITED STATES
SECURITIES AND EXCHANGE COMMISSION
Washington, D.C. 20549**

FORM 8-K

CURRENT REPORT

Pursuant to Section 13 or 15(d) of the Securities Exchange Act of 1934

Date of Report (Date of earliest event Reported): March 29, 2016

Pain Therapeutics, Inc.

(Exact Name of Registrant as Specified in Charter)

Delaware
(State or Other Jurisdiction of
Incorporation)

000-29959
(Commission File Number)

91-1911336
(I.R.S. Employer Identification Number)

7801 N Capital of Texas Highway, Suite 260, Austin, TX 78731
(Address of Principal Executive Offices) (Zip Code)

512-501-2444
(Registrant's telephone number, including area code)

Not Applicable
(Former name or former address, if changed since last report)

Check the appropriate box below if the Form 8-K filing is intended to simultaneously satisfy the filing obligation of the registrant under any of the following provisions:

- Written communications pursuant to Rule 425 under the Securities Act (17 CFR 230.425)
 - Soliciting material pursuant to Rule 14a-12 under the Exchange Act (17 CFR 240.14a-12)
 - Pre-commencement communications pursuant to Rule 14d-2(b) under the Exchange Act (17 CFR 240.14d-2(b))
 - Pre-commencement communications pursuant to Rule 13e-4(c) under the Exchange Act (17 CFR 240.13e-4(c))
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Item 8.01. Other Events.

On March 29, 2016, the Registrant issued a press release, a copy of which is attached hereto as Exhibit 99.1 and is incorporated herein by reference.

Item 9.01. Financial Statements and Exhibits.

Exhibit 99.1. Press release dated March 29, 2016

SIGNATURE

Pursuant to the requirements of the Securities Exchange Act of 1934, the registrant has duly caused this report to be signed on its behalf by the undersigned hereunto duly authorized.

Pain Therapeutics, Inc.

Date: March 29, 2016

By: /s/ Peter S. Roddy_____

Name: Peter S. Roddy

Title: Vice President & Chief Financial Officer

Pain Therapeutics Resubmits REMOXY New Drug Application to the U.S. Food and Drug Administration

REMOXY NDA Has Priority Review

AUSTIN, Texas, March 29, 2016 (GLOBE NEWSWIRE) -- Pain Therapeutics, Inc. (Nasdaq:PTIE) announced today that it has resubmitted to the U.S. Food and Drug Administration (FDA) a New Drug Application (NDA) for REMOXY®, an abuse-deterrent formulation of extended-release oxycodone (CII) capsules. Pain Therapeutics expects to be notified by the FDA of a Prescription Drug User Fee Act (PDUFA) action date within 30 days. The original REMOXY NDA has a Priority Review designation.

REMOXY is a proprietary drug developed and owned by Pain Therapeutics. The drug candidate's proposed indication is for "the management of pain severe enough to require daily, around-the-clock, long-term opioid treatment and for which alternative treatment options are inadequate".

Pain Therapeutics specifically developed REMOXY to make oxycodone difficult to abuse yet provide 12 hours of steady pain relief when used appropriately by patients. In particular, REMOXY's thick, sticky, high-viscosity formulation may deter unapproved routes of drug administration, such as injection, snorting or smoking. Pain Therapeutics owns exclusive, worldwide rights to REMOXY.

About Opioid Abuse

Opioid drugs such as oxycodone are an important treatment option for patients with severe chronic pain. However, oxycodone abuse and diversion remains a serious, persistent problem. Nearly 19,000 people died from opioid overdose in 2014, according to the National Institute on Drug Abuse. For over a decade, Pain Therapeutics has pioneered *Abuse-Deterrent Formulations* (ADFs) to help in the fight against prescription drug abuse. ADFs attempt to raise the bar on prescription drug abuse by making it difficult, longer or aversive to tamper with long-acting opioid formulations, recognizing that no drug can be made abuse-proof.

About Pain Therapeutics, Inc.

We develop proprietary drugs that offer significant improvements to patients and physicians. Our expertise consists of developing new drugs and guiding these through various regulatory and development pathways in preparation for their eventual commercialization. We generally focus our drug development efforts around disorders of the nervous system, such as chronic pain. The FDA has not yet established the safety or efficacy of our drug candidates.

NOTE: REMOXY® is a trademark of Pain Therapeutics, Inc.

Note Regarding Forward-Looking Statements: *This press release contains forward-looking statements for purposes of the Private Securities Litigation Reform Act of 1995 (the "Act"). Pain Therapeutics disclaims any intent or obligation to update these forward-looking statements, and claims the protection of the Safe Harbor for forward-looking statements contained in the Act. Examples of such statements include, but are not limited to, statements regarding a PDUFA date; the proposed indication for REMOXY; the abuse-deterrent properties of REMOXY and the potential benefits of REMOXY. Such statements are based on management's current expectations, but actual results may differ materially due to various factors. Drug development involves substantial risks and uncertainties, including, but not limited to, those risks and uncertainties relating to the FDA's acceptance to review the REMOXY NDA; difficulties or delays in the FDA's review of REMOXY; risks that the REMOXY NDA is administratively incomplete or out of compliance with current regulations around content and format or risks that the REMOXY NDA is missing key components, any of which may lead to a refusal-to-file decision by the FDA; expected adverse side effects or inadequate therapeutic efficacy; the uncertainty of patent protection for our intellectual property or trade secrets; unanticipated additional research and development, litigation and other costs; the potential for abuse-deterrent pain medications or other competing products to be developed by others; and difficulties or delays in manufacturing REMOXY. For further information regarding these and other risks related to our business, investors should consult our filings with the U.S. Securities and Exchange Commission.*

For More Information Contact:

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