
**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**

**October 11, 2010
Date of Report (Date of earliest event reported)**

JAZZ PHARMACEUTICALS, INC.

(Exact name of Registrant as specified in its charter)

Delaware
**(State or Other Jurisdiction
of Incorporation)**

001-33500
**(Commission
File No.)**

05-0563787
**(IRS Employer
Identification No.)**

3180 Porter Drive, Palo Alto, California 94304
(Address of principal executive offices, including zip code)

(650) 496-3777
(Registrant's telephone number, including area code)

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 October 11, 2010, Jazz Pharmaceuticals, Inc. issued a press release titled “Jazz Pharmaceuticals Receives FDA Complete Response Letter Regarding JZP-6 for Treatment of Fibromyalgia.” A copy of the press release is furnished as Exhibit 99.1 to this report.

Item 9.01. Financial Statements and Exhibits.

(d) Exhibits

<u>Exhibit Number</u>	<u>Description</u>
99.1	Press Release dated October 11, 2010

EXHIBIT INDEX

**Exhibit
Number**

Description

99.1

Press Release dated October 11, 2010



**JAZZ PHARMACEUTICALS RECEIVES FDA COMPLETE RESPONSE LETTER
REGARDING JZP-6 FOR TREATMENT OF FIBROMYALGIA**

— Company to host conference call on Monday, October 11 at 8:30 AM ET —

PALO ALTO, Calif., Oct. 11, 2010 — Jazz Pharmaceuticals, Inc. (Nasdaq: JAZZ) announced today that the U.S. Food and Drug Administration (FDA) has sent the company a complete response letter (CRL) regarding the company's New Drug Application (NDA) for JZP-6 (sodium oxybate) for the treatment of fibromyalgia. The CRL states that the FDA cannot approve the NDA in its present form. In the letter, the FDA discusses a number of topics, including the need for additional clinical studies, the appropriate patient population, methods for ensuring safe use, and the proposed REMS, concentration and trade name for the product.

"We have requested a meeting with FDA in order to discuss and clarify the contents of the CRL and will then evaluate our next steps for JZP-6," said Bruce Cozadd, chairman and chief executive officer of Jazz Pharmaceuticals. "We continue to believe there is a significant unmet medical need among fibromyalgia patients that could be met by JZP-6 if it were approved by FDA."

Conference Call Information

Jazz Pharmaceuticals will host an investor conference call and live audio webcast today (October 11) at 8:30 AM Eastern Time/ 5:30 AM Pacific Time to discuss this release and respond to investor questions. The live webcast may be accessed from the Investors section of the Jazz Pharmaceuticals website at www.jazzpharmaceuticals.com. Please connect to the website prior to the start of the conference call to ensure adequate time for any software downloads that may be necessary.



Investors may participate in the conference call by dialing 866-700-6067 in the U.S., or 617-213-8834 outside the U.S., and entering passcode 43104766.

An archived version of the webcast will be available for at least one week on the investors section of the Jazz Pharmaceuticals' website at www.jazzpharmaceuticals.com.

About Jazz Pharmaceuticals, Inc.

Jazz Pharmaceuticals (Nasdaq: JAZZ) is a specialty pharmaceutical company that identifies, develops and commercializes innovative treatments for important, underserved markets in neurology and psychiatry. For further information see www.jazzpharmaceuticals.com.

“Safe Harbor” Statement under the Private Securities Litigation Reform Act of 1995

This press release contains forward-looking statements, including but not limited to statements related to Jazz Pharmaceuticals' JZP-6 (sodium oxybate) product candidate, including statements related to Jazz Pharmaceuticals' plans to meet with the FDA, the outcome of any such meeting and all future regulatory matters, including its future approval and the future development of JZP-6. These forward-looking statements are based on the company's current expectations and inherently involve significant risks and uncertainties. Jazz Pharmaceuticals' actual results and the timing of events could differ materially from those anticipated in such forward-looking statements as a result of these risks and uncertainties, which include, without limitation, risks related to the uncertain and time-consuming regulatory approval process for JZP-6, risks related to potential further development of JZP-6, and other risks detailed from time-to-time under the caption “Risk Factors” and elsewhere in Jazz Pharmaceuticals' Securities and Exchange Commission filings and reports, including in its Quarterly Report on Form 10-Q for the quarter ended June 30, 2010 filed by Jazz Pharmaceuticals with the Securities and Exchange Commission on August, 11, 2010. Jazz Pharmaceuticals undertakes no duty or obligation to update any forward-looking statements contained in this release as a result of new information, future events or changes in its expectations.

Contact

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