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

January 7, 2008
Date of Report (Date of earliest event reported)

JAZZ PHARMACEUTICALS, INC.

(Exact name of Registrant as specified in its charter)

Delaware	**001-33500**	**05-0563787**
(State or Other Jurisdiction of Incorporation)	**(Commission File No.)**	**(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))

Item 8.01 Other Events.

On January 7, 2008, Jazz Pharmaceuticals, Inc. announced that it has received orphan drug designation from the U.S. Food and Drug Administration for its JZP-8 product candidate that is being developed for the treatment of recurrent acute repetitive seizures. A copy of the press release is attached as Exhibit 99.1 to this current report and is incorporated by reference herein.

Item 9.01 Financial Statements and Exhibits.

(d) Exhibits

Exhibit Number	Description
99.1	Press Release dated January 7, 2008

SIGNATURES

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.

JAZZ PHARMACEUTICALS, INC.

By: /s/ Carol A. Gamble
Carol A. Gamble
Senior Vice President, General Counsel
and Corporate Secretary

Date: January 8, 2008

EXHIBIT INDEX

Exhibit Number	Description
99.1	Press Release dated January 7, 2008

Exhibit 99.1

JAZZ PHARMACEUTICALS, INC. ANNOUNCES RECEIPT OF FDA ORPHAN DRUG DESIGNATION FOR RECURRENT ACUTE REPETITIVE SEIZURES PRODUCT CANDIDATE

PALO ALTO, Calif. January 7, 2008 – Jazz Pharmaceuticals, Inc. (Nasdaq: JAZZ) announced today that it has received orphan drug designation from the U.S. Food and Drug Administration (FDA) for its JZP-8 product candidate for the treatment of recurrent acute repetitive seizures. In December 2007, Jazz Pharmaceuticals dosed the first patient in a Phase II clinical trial of JZP-8.

JZP-8 is a novel drug delivery formulation incorporating clonazepam, a widely prescribed benzodiazepine. The product candidate is designed to be a fast-acting intranasal spray for the treatment of recurrent acute repetitive seizures in patients with epilepsy. Jazz Pharmaceuticals previously completed development activities to select the active pharmaceutical ingredient for the product, to determine its formulation, and to assess its safety and tolerability in early stage studies.

“JZP-8 is one of the first development product candidates from our internal product identification and development program to enter Phase II,” said Samuel R. Saks, M.D., Chief Executive Officer of Jazz Pharmaceuticals. “Dosing of the first patient in our Phase II clinical trial and receipt of orphan drug designation are important milestones for this program.”

About Recurrent Acute Repetitive Seizures

Recurrent acute recurrent repetitive seizures (ARS) are bouts of acute seizure activity occurring within a 24-hour period in adults and a 12-hour period in children. ARS, also called seizure clusters, occurs in a small subset of patients with epilepsy who regularly experience breakthrough seizures in flurries or clusters despite treatment with a regimen of anti-epileptic drugs. ARS is a rare disorder, with an estimated prevalence in the United States of approximately 90,000 to 180,000 people.

About Orphan Drug Designation

Orphan drug designation is granted by the FDA to encourage the development of treatments for diseases or conditions that affect fewer than 200,000 patients in the United States. Drug products that receive orphan drug designation obtain seven years of marketing exclusivity from the date of FDA marketing approval, as well as tax credits for clinical trial costs and waivers of marketing application filing fees.

About Jazz Pharmaceuticals, Inc.

Jazz Pharmaceuticals, Inc. is a specialty pharmaceutical company focused on identifying, developing and commercializing innovative products to meet unmet medical needs in neurology and psychiatry. For further information, please visit 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 the development program for the JZP-8 product candidate and the company’s internal product concept generation program. These forward-looking statements 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, whether or when clinical studies of JZP-8 will be completed or will be successful, and whether or when the product candidate will be approved or launched. These and other risk factors are discussed under “Risk Factors,” in the Quarterly Report on Form 10-Q for the quarter ended September 30, 2007 filed by Jazz Pharmaceuticals with the Securities and Exchange Commission on November 9, 2007. 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.

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