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): June 11, 2010

QUESTCOR PHARMACEUTICALS, INC.

(Exact Name of Registrant as Specified in Charter)

California	**001-14758**	**33-0476164**
(State or Other Jurisdiction of Incorporation)	(Commission File Number)	(I.R.S. Employer Identification No.)

3260 Whipple Road, Union City, California	**94587**
(Address of Principal Executive Offices)	(Zip Code)

Registrant's telephone number, including area code: **(510) 400-0700**

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 (see General Instruction A.2. below):

- o Written communications pursuant to Rule 425 under the Securities Act (17 CFR 230.425)
- o Soliciting material pursuant to Rule 14a-12 under the Exchange Act (17 CFR 240.14a-12)
- o Pre-commencement communications pursuant to Rule 14d-2(b) under the Exchange Act (17 CFR 240.14d-2(b))
- o 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 June 11, 2010, Questcor Pharmaceuticals, Inc. (the "Company") was informed by the U.S. Food & Drug Administration ("FDA") that the PDUFA date for Questcor's supplemental new drug application (sNDA) to approve Acthar for the treatment of infantile spasms (IS) has been extended to September 11, 2010. The FDA extended the PDUFA date in order to review information regarding labeling and potential post-approval commitments that they solicited from Questcor. This follows the May 6, 2010 votes by the Advisory Committee to the Division of Peripheral and Central Nervous System Drugs of the FDA which indicated support for approval of this new Acthar indication.

Investors should note that there can be no assurance that the FDA will approve this sNDA by September 11, 2010, or thereafter.

A copy of the press release announcing the FDA's extension of the PDUFA date is filed as Exhibit 99.1 hereto and incorporated by reference herein.

Item 9.01. Financial Statements and Exhibits.

(c) Exhibits.

Exhibit No.	Description
99.1	Questcor Pharmaceuticals, Inc. press release dated June 11, 2010.

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.

Date: June 14, 2010

QUESTCOR PHARMACEUTICALS, INC.

By: /s/ Gary M. Sawka
Gary M. Sawka
Senior Vice-President, Finance and Chief Financial Officer

EXHIBIT INDEX

Exhibit No.	Description
99.1	Questcor Pharmaceuticals, Inc. press release dated June 11, 2010.

Exhibit 99.1

QUESTCOR RECEIVES NOTIFICATION OF PDUFA DATE EXTENSION TO SEPTEMBER 11, 2010

Union City, CA — June 11, 2010 — Questcor Pharmaceuticals, Inc. (NASDAQ: QCOR) today announced that it has received notification from the U.S. Food & Drug Administration (FDA) that the PDUFA date for Questcor's supplemental new drug application (sNDA) to approve Acthar for the treatment of infantile spasms (IS) has been extended to September 11, 2010. The FDA extended the PDUFA date in order to review information regarding labeling and potential post-approval commitments that they solicited from Questcor. This follows the May 6, 2010 votes by the Advisory Committee to the Division of Peripheral and Central Nervous System Drugs of the FDA which indicated support for approval of this new Acthar indication.

"We look forward to working with the FDA over the next few months in order to finalize these critical elements," said Don M. Bailey, President and CEO of Questcor.

Investors should note that there can be no assurance that the FDA will approve this sNDA by September 11, 2010, or thereafter.

About Questcor

Questcor Pharmaceuticals, Inc. is a pharmaceutical company that markets H.P. Acthar® Gel (repository corticotropin injection). H.P. Acthar Gel ("Acthar") is an injectable drug that is approved for the treatment of certain disorders with an inflammatory component, including the treatment of exacerbations associated with multiple sclerosis ("MS") and to induce a diuresis or a remission of proteinuria in the nephrotic syndrome without uremia of the idiopathic type or that is due to lupus erythamatosus. In addition, Acthar is not indicated for, but is used in treating patients with infantile spasms ("IS"), a rare form of refractory childhood epilepsy, and opsoclonus myoclonus syndrome, a rare autoimmune-related childhood neurological disorder. The Company also markets Doral® (quazepam), which is indicated for the treatment of insomnia characterized by difficulty in falling asleep, frequent nocturnal awakenings, and/or early morning awakenings. For more information, please visit www.questcor.com.

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