
**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): September 19, 2012

QUESTCOR PHARMACEUTICALS, INC.

(Exact Name of Registrant as Specified in Charter)

California
(State or Other Jurisdiction
of Incorporation)

001-14758
(Commission
File Number)

33-0476164
(I.R.S. Employer
Identification No.)

1300 Kellogg Drive, Suite D, Anaheim, California
(Address of Principal Executive Offices)

92807
(Zip Code)

Registrant's telephone number, including area code: (714) 786-4200

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 7.01. Regulation FD Disclosure.

On September 19, 2012, Questcor Pharmaceuticals, Inc. (the “Company”) issued a press release regarding a policy bulletin issued in respect of the Company’s primary product H.P. Acthar® Gel (repository corticotropin injection). A copy of the Company’s press release is attached hereto as Exhibit 99.1 and incorporated herein by reference.

In accordance with General Instruction B.2. of Form 8-K, the information in Item 7.01 of this Current Report on Form 8-K, including Exhibit 99.1, shall not be deemed “filed” for purposes of Section 18 of the Securities Exchange Act of 1934, as amended (the “Exchange Act”), or otherwise subject to the liability of that section, nor shall it be deemed incorporated by reference in any filing under the Securities Act of 1933, as amended, or the Exchange Act, except as expressly set forth by specific reference in such a filing.

Item 9.01. Financial Statements and Exhibits.

(d) Exhibits.

<u>Exhibit No.</u>	<u>Description</u>
99.1	Questcor Pharmaceuticals, Inc. Press Release dated September 19, 2012.

SIGNATURES

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

Date: September 19, 2012

QUESTCOR PHARMACEUTICALS, INC.

By: /s/ Michael H. Mulroy

Michael H. Mulroy
Senior Vice President, Chief Financial Officer,
and General Counsel

EXHIBIT INDEX

<u>Exhibit No.</u>	<u>Description</u>
99.1	Questcor Pharmaceuticals, Inc. Press Release dated September 19, 2012.



Questcor Comments on Insurance Policy Bulletin

ANAHEIM, Calif., September 19, 2012 — Questcor Pharmaceuticals, Inc. (NASDAQ: QCOR) today commented on information regarding an insurance policy bulletin issued with respect to the Company's primary product H.P. Acthar® Gel (repository corticotropin injection) ("Acthar").

The Company is continuing to review the Clinical Policy Bulletin related to Acthar from Aetna Inc. ("Aetna"). Currently, the Company does not believe that the bulletin represents a material change in insurance coverage for Acthar by Aetna. During 2012, Aetna has accounted for approximately 5% of the Company's shipped prescriptions for Acthar. Based on its current assessment of the Clinical Policy Bulletin, the Company does not believe that the bulletin will have a material impact on the Company's results of operations.

About Questcor

Questcor Pharmaceuticals, Inc. is a biopharmaceutical company focused on the treatment of patients with serious, difficult-to-treat autoimmune and inflammatory disorders. Questcor's primary product is H.P. Acthar® Gel (repository corticotropin injection), an injectable drug that is approved by the FDA for the treatment of 19 indications. Of these 19 indications, Questcor currently generates substantially all of its net sales from three indications: the treatment of proteinuria in idiopathic types of nephrotic syndrome, the treatment of acute exacerbations of multiple sclerosis in adults, and the treatment of infantile spasms in children under two years of age. With respect to nephrotic syndrome, the FDA has approved Acthar to "induce a diuresis or a remission of proteinuria in the nephrotic syndrome without uremia of the idiopathic type or that due to lupus erythematosus." Questcor is also exploring the possibility of developing markets for other on-label indications and the possibility of pursuing FDA approval of additional indications not currently on the Acthar label where there is high unmet medical need. For more information about Questcor, please visit www.questcor.com.



Note: Except for the historical information contained herein, this press release contains forward-looking statements that have been made pursuant to the Private Securities Litigation Reform Act of 1995. These statements relate to future events or our future financial performance. In some cases, you can identify forward-looking statements by terminology such as “believes,” “continue,” “could,” “estimates,” “expects,” “growth,” “may,” “plans,” “potential,” “should,” “substantial” or “will” or the negative of such terms and other comparable terminology. These statements are only predictions. Actual events or results may differ materially. Factors that could cause or contribute to such differences include, but are not limited to, the following:

- Our reliance on Acthar for substantially all of our net sales and profits;
- Our ability to continue to generate revenue from sales of Acthar to treat on-label indications associated with NS, and our ability to develop other therapeutic uses for Acthar;
- Our ability to receive high reimbursement levels from third party payers;
- An increase in the proportion of our Acthar unit sales comprised of Medicaid-eligible patients and government entities; and
- Other risks discussed in Questcor’s annual report on Form 10-K for the year ended December 31, 2011 as filed with the Securities and Exchange Commission, or SEC, on February 22, 2012, and other documents filed with the SEC.

The risk factors and other information contained in these documents should be considered in evaluating Questcor’s prospects and future financial performance.

Questcor undertakes no obligation to publicly release the result of any revisions to these forward-looking statements, which may be made to reflect events or circumstances after the date of this release.

For more information, please visit www.questcor.com or www.acthar.com.

CONTACT INFORMATION:

Questcor Pharmaceuticals, Inc.

Don Bailey

714-786-4210

dbailey@Questcor.com

EVC Group

Investors

Gregory Gin/Bob Jones/

646-445-4801/646-445-5447

Doug Sherk

415-568-4887

Media

Janine McCargo

646-688-0425