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

FORM 6-K

REPORT OF FOREIGN PRIVATE ISSUER
PURSUANT TO RULE 13A-16 OR 15D-16 UNDER THE
SECURITIES EXCHANGE ACT OF 1934

Dated February 11, 2016

Commission File Number 001-36421

AURINIA PHARMACEUTICALS INC.
(Exact name of Registrant as specified in its charter)

N/A
(Translation of Registrant's Name)

#1203-4464 Markham Street
Victoria, British Columbia
V8Z7X8
(250) 708-4272
(Address and telephone number of registrant's principle executive offices)

Indicate by check mark whether the registrant files or will file annual reports under cover of Form 20-F or Form 40-F:

Form 20-F ☐ Form 40-F ☒

Indicate by check mark if the registrant is submitting the Form 6-K in paper as permitted by Regulation S-T Rule 101(b)(1): ☐

Indicate by check mark if the registrant is submitting the Form 6-K in paper as permitted by Regulation S-T Rule 101(b)(7): ☐

Indicate by check mark whether by furnishing the information contained in this Form, the registrant is also thereby furnishing the information to the Commission pursuant to Rule 12g3-2(b) under the Securities Exchange Act of 1934.

Yes ☐ No ☒

If "Yes" is marked, indicate below the file number assigned to the registrant in connection with Rule 12g3-2(b): Not applicable.

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, thereunto duly authorized.

Dated: February 11, 2016

Aurinia Pharmaceuticals Inc.

By: /s/ Michael R. Martin

Name: Michael R. Martin

Title: Chief Operating Officer

EXHIBIT INDEX

Exhibit	Description of Exhibit
99.1	News Release – Aurinia to Host Conference Call & Webcast to Review Results from the AURION Study

Exhibit 99.1

Aurinia to Host Conference Call & Webcast to Review Results from the AURION Study

Conference Call and Webcast with Slides, Tuesday February 16th at 4.30pm Eastern Time

VICTORIA, British Columbia--(BUSINESS WIRE)--February 11, 2016--**Aurinia Pharmaceuticals Inc**. (NASDAQ:AUPH / TSX:AUP) ("Aurinia" or the "Company") will host a conference call and live webcast to discuss detailed results from its open label, single arm exploratory AURION (***A***urinia early ***U***rinary protein ***R***educt***ion*** Predicts Response) study on Tuesday, February 16 at 4:30pm Eastern Standard Time. Participating in the call will be Stephen Zaruby, CEO, Dr. Neil Solomons, Chief Medical Officer, and Michael Martin, Chief Operating Officer.

Tuesday, February 16, 2016 @ 4:30pm Eastern/1:30pm Pacific
Domestic: 888-329-8862
International: 719-785-1753
Conference ID: 3145908
Webcast with slides: http://public.viavid.com/index.php?id=118297

Replays, through March 1, 2016
Domestic: 877-870-5176
International: 858-384-5517
Conference ID: 3145908

About Aurinia

Aurinia is a clinical stage pharmaceutical company focused on the global nephrology market. The fully-enrolled 265 patient Phase 2b AURA-LV clinical trial is evaluating the efficacy of its lead drug, voclosporin, as a treatment for active LN. LN is an inflammation of the kidneys, that if inadequately treated can lead to end-stage renal disease, making LN a serious and potentially life-threatening condition.

Voclosporin is a novel and potentially best-in-class calcineurin inhibitor ("CNI") with extensive clinical data in over 2,000 patients in other indications. Voclosporin is made by a modification of a single amino acid of the cyclosporine molecule (a CNI approved for use in transplant patients since 1983). This modification results in a more predictable pharmacokinetic and pharmacodynamic relationship, an increase in potency vs. cyclosporine, an altered metabolic profile, and potential for flat dosing.

About AURION:

The AURION study or "Aurinia Early Urinary Protein Reduction Predicts Response Study" is an open label, exploratory study being conducted in multiple sites in Malaysia to assess the short term predictors of response using voclosporin (23.7mg) in combination with mycophenolate mofetil and oral corticosteroids in patients with active lupus nephritis. This study will examine biomarkers of disease activity at 8 weeks and their ability to predict response at 24 and 48 weeks.

We seek Safe Harbor.

CONTACT:
For More Information:
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