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 December 9, 2019

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 principal executive offices)

Indicate by check mark whether the registrant files or will file annual reports under cover of Form20-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 ☐

This Form 6-K is hereby filed and incorporated by reference into the Registrant's Registration Statement on FormF-10 (File No. 333-222413).

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: December 9, 2019

Aurinia Pharmaceuticals Inc.

By: /s/ Peter S. Greenleaf
Name: Peter S. Greenleaf
Title: Chief Executive Officer

EXHIBIT INDEX

Exhibit	Description of Exhibit
99.1	News Release - Aurinia Announces Public Offering of Common Shares
99.2	Material Change Report, dated December 9, 2019 - AURORA Clinical Trial Results

Exhibit 99.1 included with this report on Form 6-K is hereby incorporated by reference as an exhibit to the Registrant's Registration Statement on Form F-10 (File No. 333-222413), as amended or supplemented.

Aurinia Announces Public Offering of Common Shares

VICTORIA, British Columbia—(BUSINESS WIRE)—December 9, 2019—Aurinia Pharmaceuticals Inc. (NASDAQ:AUPH) (TSX:AUP) ("Aurinia" or the "Company"), a late- stage clinical biopharmaceutical company focused on advancing voclosporin in multiple indications, today announced that it has commenced a registered underwritten public offering of US$150,000,000 of its common shares (the "Offering").

Jefferies LLC and SVB Leerink LLC are acting as joint book-running managers for the Offering. H.C. Wainwright & Co., LLC, Oppenheimer & Co. Inc. and Bloom Burton Securities Inc. are acting as co-managers for the Offering.

The Company will grant the underwriters an option exercisable, in whole or in part, in the sole discretion of the underwriters, to purchase up to an aggregate of US$22,500,000 of additional shares, for a period of up to 30 days. The Offering is subject to market conditions, and there can be no assurance as to whether or when the Offering may be completed, or as to the actual size or terms of the Offering.

The Company intends to use the net proceeds of the Offering forpre-commercialization and launch activities, as well as working capital and general corporate purposes.

The Offering is subject to customary closing conditions, including NASDAQ and TSX approvals. For the purposes of the TSX approval, the Company intends to rely on the exemption set forth in Section 602.1 of the TSX Company Manual, which provides that the TSX will not apply its standards to certain transactions involving eligible interlisted issuers on a recognized exchange, such as NASDAQ.

The Offering is being made pursuant to a U.S. registration statement on FormF-10, declared effective by the United States Securities and Exchange Commission (the "SEC") on March 29, 2018 (the "Registration Statement"), and the Company's existing Canadian short form base shelf prospectus (the "Base Shelf Prospectus") dated March 26, 2018. The prospectus supplements relating to the Offering (together with the Base Shelf Prospectus and the Registration Statement, the "Offering Documents") will be filed with the securities commissions in the provinces of British Columbia, Alberta and Ontario in Canada, and with the SEC in the United States. The Offering Documents will contain important detailed information about the securities being offered. Before you invest, you should read the Offering Documents and the other documents the Company has filed for more complete information about the Company and the Offering. Copies of the Offering Documents will be available for free by visiting the Company's profiles on the SEDAR website maintained by the Canadian Securities Administrators at www.sedar.com or the SEC's website at www.sec.gov, as applicable. Alternatively, copies of the prospectus supplement will be available upon request by contacting Jefferies LLC, Attention: Equity Syndicate Prospectus Department, 520 Madison Avenue, 2nd Floor, New York, NY 10022; by phone at (877) 821-7388; or by e-mail at Prospectus_Department@Jefferies.com; or SVB Leerink LLC, Attention: Syndicate Department, One Federal Street, 37th Floor, Boston, MA 02110, by telephone at 1-800-808-7525, ext. 6132, or by email at syndicate@svbleerink.com.

This press release does not constitute an offer to sell or the solicitation of an offer to buy securities, nor will there be any sale of the securities in any jurisdiction in which such offer, solicitation or sale would be unlawful prior to the registration or qualification under the securities laws of any such jurisdiction.

ABOUT AURINIA

Aurinia Pharmaceuticals is a late clinical-stage biopharmaceutical company focused on developing and commercializing therapies to treat targeted patient populations that are impacted by serious diseases with a high unmet medical need. The Company is currently developing an investigational drug, for the treatment of Lupus Nephritis, Focal Segmental Glomerulosclerosis and Dry Eye Syndrome. The Company's head office is in Victoria, British Columbia and focuses its development efforts globally.

Investors & Media:

Glenn Schulman, PharmD, MPH
SVP, Corporate Communications & IR
gschulman@auriniapharma.com

FORM 51-102F3
Material Change Report

Item 1 **Name and Address of Company**

Aurinia Pharmaceuticals Inc. (the "**Company**")
#1203-4464 Markham Street
Victoria, BC V8Z 7X8
Canada

Item 2 **Date of Material Change**

December 4, 2019

Item 3 **News Release**

A news release was issued and disseminated by the Company throughBusiness News Wire on December 4, 2019.

Item 4 **Summary of Material Change**

The Company announced positive efficacy and safety results from its pivotal AURORA Phase 3 trial of voclosporin, in combination with mycophenolate ("**MMF**") and low-dose corticosteroids, in the treatment of lupus nephritis ("**LN**").

Item 5 **Full Description of Material Change**

The Company announced positive efficacy and safety results from its pivotal AURORA Phase 3 trial of voclosporin, in combination with MMF and low-dose corticosteroids, in the treatment of LN.

This global study, in which 357 patients with active LN were enrolled, met its primary endpoint of achieving renal response at 52 weeks, demonstrating response rates of 40.8% for voclosporin vs. 22.5% for the control (OR 2.65; $p < 0.001$). Additionally, all pre-specified hierarchical secondary endpoints achieved statistical significance in favor of voclosporin, which included renal response at 24 weeks, partial renal response at 24 and 52 weeks, time to achieve urinary protein-to-creatinine ratio ("**UPCR**") £ 0.5, and time to 50% reduction in UPCR. The robustness of the data was also supported by all pre-specified subgroup analyses (age, sex, race, biopsy class, region, and prior MMF use) favoring voclosporin.

	Measure	Result	Odds Ratio [95% CI]	p-value
Primary Endpoint	Renal Response at 52 weeks	Voclosporin 40.8% Control 22.5%	2.65 [1.64, 4.27]	p < 0.001
Secondary Endpoints	Renal Response at 24 weeks	Voclosporin 32.4% Control 19.7%	2.23 [1.34, 3.72]	p = 0.002
	Partial Renal Response at 24 weeks	Voclosporin 70.4% Control 50.0%	2.43 [1.56, 3.79]	p < 0.001
	Partial Renal Response at 52 weeks	Voclosporin 69.8% Control 51.7%	2.26 [1.45, 3.51]	p < 0.001
	Time to UPCR £ 0.5	Voclosporin faster than Control	2.02 [1.51, 2.70] Hazard Ratio	p < 0.001
	Time to 50% reduction in UPCR	Voclosporin faster than Control	2.05 [1.62, 2.60] Hazard Ratio	p < 0.001

Voclosporin was generally well tolerated with no unexpected safety signals. Serious adverse events (“**SAEs**”) were reported in 20.8% of voclosporin patients vs. 21.3% in the control arm. Infection was the most commonly reported SAE with 10.1% of voclosporin patients versus 11.2% of patients in the control arm. Overall mortality in the trial was low, with six deaths observed; one in the voclosporin arm and five in the control group. None of the deaths were determined to be treatment related. Additionally, the voclosporin arm showed no significant decrease at week 52 in estimated glomerular filtration rate or increase in blood pressure, lipids or glucose, which are common adverse events associated with legacy calcineurin inhibitors (“**CNIs**”).

Voclosporin was granted Fast Track designation by the Food and Drug Administration of the United States Government (“**FDA**”) in 2016. The Company expects to hold a pre-New Drug Application (“**NDA**”) meeting with the FDA in the first quarter of 2020, and submit a NDA to the FDA in the first half of 2020.

Item 5.2 Disclosure of Restructuring Transactions

Not applicable.

Item 6 Reliance on subsection 7.1(2) of National Instrument 51-102

Not applicable.

Item 7 Omitted Information

No significant facts remain confidential in, and no information has been omitted from, this report.

Item 8 Executive Officer

For further information, please contact:

Mr. Michael R. Martin, Chief Operating Officer
250-708-4272
mmartin@auriniapharma.com

Item 9 Date of Report

December 9, 2019

Cautionary Note Regarding Forward-looking Statements

Certain statements made in this press release may constitute forward-looking information within the meaning of applicable Canadian securities law and forward-looking statements within the meaning of applicable United States securities law. These forward-looking statements or information include but are not limited to statements or

information with respect to the anticipated pre-NDA meeting with the FDA in the first quarter of 2020 and the anticipated NDA filing to the FDA during the first half of 2020. It is possible that such results or conclusions may change based on further analyses of these data. Words such as "anticipate", "will", "believe", "estimate", "expect", "intend", "target", "plan", "goals", "objectives", "may" and other similar words and expressions, identify forward-looking statements. We have made numerous assumptions about the forward-looking statements and information contained herein, including among other things, assumptions about: the market value for the LN, dry eye syndrome ("DES") and focal segmental glomerulosclerosis ("FSGS") programs; that another company will not create a substantial competitive product for Aurinia's LN, DES and FSGS business without violating Aurinia's intellectual property rights; the burn rate of Aurinia's cash for operations; the costs and expenses associated with Aurinia's clinical trials; the planned studies achieving positive results; Aurinia being able to extend and protect its patents on terms acceptable to Aurinia; and the size of the LN, DES or FSGS markets; Aurinia will be able to obtain all necessary regulatory approvals for commercialization of voclosporin for use in LN on terms that are acceptable to it and that are commercially viable; and that Aurinia's intellectual property rights are valid and do not infringe the intellectual property rights of other parties. Even though the management of Aurinia believes that the assumptions made, and the expectations represented by such statements or information are reasonable, there can be no assurance that the forward-looking information will prove to be accurate.

Forward-looking information by their nature are based on assumptions and involve known and unknown risks, uncertainties and other factors which may cause the actual results, performance or achievements of Aurinia to be materially different from any future results, performance or achievements expressed or implied by such forward-looking information. Should one or more of these risks and uncertainties materialize, or should underlying assumptions prove incorrect, actual results may vary materially from those described in forward-looking statements or information. Such risks, uncertainties and other factors include, among others, the following: difficulties, delays, or failures we may experience in the conduct of our clinical trial; difficulties we may experience in completing the development and commercialization of voclosporin; the market for the LN, DES and FSGS business may not be as estimated; Aurinia may have to pay unanticipated expenses; estimated costs for clinical trials may be underestimated, resulting in Aurinia having to make additional expenditures to achieve its current goals; Aurinia not being able to extend or fully protect its patent portfolio for voclosporin; competitors may arise with similar products; Aurinia may not be able to obtain necessary regulatory approvals for commercialization of voclosporin in a timely fashion, or at all; and Aurinia may not be able to obtain sufficient supply to meet commercial demand for voclosporin in a timely fashion. Although we have attempted to identify factors that would cause actual actions, events or results to differ materially from those described in forward-looking statements and information, there may be other factors that cause actual results, performances, achievements or events to not be as anticipated, estimated or intended. Also, many of the factors are beyond our control. There can be no assurance that forward-looking statements or information will prove to be accurate, as actual results and future events could differ materially from those anticipated in such statements. Accordingly, you should not place undue reliance on forward-looking statements or information.

Except as required by law, Aurinia will not update forward-looking information. All forward-looking information contained in this press release is qualified by this cautionary statement. Additional information related to Aurinia, including a detailed list of the risks and uncertainties affecting Aurinia and its business can be found in Aurinia's most recent Annual Information Form available by accessing the Canadian Securities Administrators' System for Electronic Document Analysis and Retrieval (SEDAR) website at www.sedar.com or the U.S. Securities and Exchange Commission's Electronic Document Gathering and Retrieval System (EDGAR) website at www.sec.gov/edgar.