

ARQULE INC  
Form 8-K  
April 17, 2018

**UNITED STATES**

**SECURITIES AND EXCHANGE COMMISSION**

**Washington, DC 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): April 16, 2018

**ARQULE, INC.**

(Exact name of registrant as specified in its charter)

**Delaware**

(State or other jurisdiction  
of incorporation)

**000-21429**

(Commission File Number)

**04-3221586**

(I.R.S. Employer

Identification No.)

**One Wall Street**

**Burlington, MA 01803**

(Address of principal executive offices) (Zip Code)

**(781) 994-0300**

(Registrant's telephone number, including area code)

**N/A**

(Former name or former address, if changed since last report)

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):

☐ 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))

Indicate by check mark whether the registrant is an emerging growth company as defined in Rule 405 of the Securities Act of 1933 (§230.405 of this chapter) or Rule 12b-2 of the Securities Exchange Act of 1934 (§240.12b-2 of this chapter).

☐ Emerging growth company

If an emerging growth company, indicate by check mark if the registrant has elected not to use the extended transition period for complying with any new or revised financial accounting standards provided pursuant to Section 13(a) of the Exchange Act. ☐

**Item 1.01. Entry into a Material Definitive Agreement.**

On April 17, 2018, ArQule, Inc. (“ArQule” or the “Registrant”) and Basilea Pharmaceutica International Limited (“Basilea”) entered into a License Agreement (the “Agreement”) pursuant to which ArQule granted Basilea an exclusive license to develop, manufacture and commercialize its FGFR inhibitor, derazantinib (ARQ 087), in the United States, EU, Japan and the rest of the world, excluding the People’s Republic of China, Hong Kong, Macau and Taiwan.

Under the terms of the agreement, ArQule will receive an upfront payment of \$10 million and is eligible for up to \$326 million in regulatory and commercial milestones. ArQule is also entitled to receive staggered single-digit to double-digit royalties on net sales upon commercialization. Basilea will be responsible for all costs and expenses of development, manufacture and commercialization in its territory. Under certain circumstances, ArQule may have the opportunity to promote derazantinib in the US directly.

A copy of the Agreement will be filed as an exhibit to the Registrant’s Quarterly Report on Form 10-Q for the second quarter ended June 30, 2018. A copy of the Registrant’s April 17, 2018 press release announcing the transaction is filed as exhibit 99.1 to this report and is incorporated herein by reference

**Item 9.01 Financial Statements and Exhibits.**

**(d) Exhibits.**

| <b>Exhibit<br/>No.</b> | <b>Description</b> |
|------------------------|--------------------|
|------------------------|--------------------|

|             |   |
|-------------|---|
| <u>99.1</u> | <u>Press release dated April 17, 2018</u> |
|-------------|---|

**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.

ArQule, Inc. (Registrant)

/s/ Peter S. Lawrence  
Peter S. Lawrence  
President and Chief Operating Officer

Date: April 17, 2018