ARRAY BIOPHARMA INC
Form 8-K
July 26, 2018

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): July 26, 2018

Array BioPharma Inc.
(Exact name of registrant as specified in its charter)

Delaware	001-16633	84-1460811
(State or other jurisdiction of incorporation)	(Commission File Number)	(I.R.S. Employer Identification No.)

3200 Walnut Street, Boulder, Colorado 80301
(Address of principal executive offices, including Zip Code)

303 381-6600
(Registrant’s telephone number, including area code)

(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:

¨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. ¨

In this report, “Array BioPharma,” “Array,” “we,” “us” and “our” refer to Array BioPharma Inc., unless the context otherwise provides.

Item 7.01 Regulation FD Disclosure.

On July 26, 2018, AstraZeneca and MSD reported that the selumetinib ASTRA trial in differentiated thyroid cancer (DTC) did not meet its primary endpoint. The ASTRA trial is a randomized, double-blind, Phase III trial in high-risk DTC. Trial results demonstrated that treatment with a short course of selumetinib and single dose adjuvant radioactive iodine therapy (RAI) did not meet its primary endpoint of improvement in complete remission (CR) rate compared to placebo. The full data will be presented at a forthcoming medical meeting.

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: July 26, 2018 Array BioPharma Inc.

By: /s/ JASON HADDOCK
Jason Haddock
Chief Financial Officer

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