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): June 16, 2022

Vaxart, Inc.

(Exact name of registrant as specified in its charter)

Delaware	001-35285	59-1212264
(State or other jurisdiction of incorporation)	(Commission File Number	er) (IRS Employer Identification No.)
170 Harbor Way, Suite 300, South San Francisco, California		94080
(Address of principal executive offices)		(Zip Code)
Registrant's tel	ephone number, including area co	de: (650) 550-3500
(Former Name	Not Applicable or Former Address, if Changed S	Since Last Report)
Check the appropriate box below if the Form 8-K filing is following provisions:	s intended to simultaneously satisfy	the filing obligation of the registrant under any of the
☐ Written communications pursuant to Rule 425 under	r the Securities Act (17 CFR 230.42	5)
□ Soliciting material pursuant to Rule 14a-12 under the Exchange Act (17 CFR 240.14a-12)		
☐ Pre-commencement communications pursuant to Ru	ule 14d-2(b) under the Exchange Ac	t (17 CFR 240.14d-2(b))
☐ Pre-commencement communications pursuant to Re	ule 13e-4(c) under the Exchange Ac	t (17 CFR 240.13e-4(c))
Securities registered pursuant to Section 12(b) of the Act:		
Title of each class	Trading symbol	Name of each exchange on which registered
Common Stock, \$0.0001 par value	VXRT	The Nasdaq Capital Market
Indicate by check mark whether the registrant is an emerg chapter) or Rule 12b-2 of the Securities Exchange Act of		Rule 405 of the Securities Act of 1933 (§230.405 of this
	1 /	Emerging Growth Company \square
If an emerging growth company, indicate by check mark to revised financial accounting standards provided pursua		e the extended transition period for complying with any new Act. \Box
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Item 8.01. Other Events.

On June 16, 2022, Vaxart, Inc. (the "Company") issued a press release announcing that it will hold a live question-and-answer session via webcast with investors on June 22, 2022, at 1:00 p.m. Eastern Time, to provide an overview of the Company's oral vaccine programs and to discuss the proposals being submitted at the Company's annual meeting of stockholders, which is now being held on July 6, 2022, following its adjournment. The close of business on April 11, 2022, will continue to be the record date for the determination of stockholders of the Company entitled to vote at the annual meeting. A copy of the press release is furnished as Exhibit 99.1 to this current report on Form 8-K.

The information in this current report on Form 8-K, including Exhibit 99.1 attached hereto, is being furnished and shall not be deemed "filed" for purposes of Section 18 of the Securities Exchange Act of 1934, as amended, or subject to the liabilities of that section or Sections 11 and 12(a)(2) of the Securities Act of 1933, as amended. The information contained in this current report and in the accompanying Exhibit 99.1 shall not be deemed incorporated by reference into any filing with the U.S. Securities and Exchange Commission made by the Company, whether made before or after the date hereof regardless of any general incorporation language in such filing.

Item 9.01. Financial Statements and Exhibits.

- (d) Exhibits.
 - 99.1 Press release issued by Vaxart, Inc. on June 16, 2022.
 - 104 Cover Page Interactive Data File (embedded within the Inline XBRL document).

SIGNATURES

Pursuant to the requirements of the Securities Exchange Act of 1934, as amended, the registrant has duly caused this report to be signed on its behalf by the undersigned hereunto duly authorized.

Vaxart, Inc.

Dated: June 16, 2022

By: /s/ Andrei Floroiu

Andrei Floroiu

Chief Executive Officer

Vaxart to Host Investor Q&A Webcast

Management to discuss progress on its programs, annual stockholders' meeting proposals

Webcast to be held on June 22 at 1:00 p.m. ET

SOUTH SAN FRANCISCO, Calif., June 16, 2022 – Vaxart, Inc. (Nasdaq: VXRT) today announced that it will host a question-and-answer webcast with investors and analysts on Wednesday, June 22, 2022, at 1:00 p.m. Eastern Time. During the webcast, Andrei Floroiu, President and Chief Executive Officer, Dr. Sean Tucker, SVP and Chief Scientific Officer, and Dr. James Cummings, Chief Medical Officer, will provide an overview of the Company's oral vaccine programs.

The Company will take written questions from investors and analysts. Please visit the webcast link to RSVP and submit written questions. Questions may also be submitted in advance to ir@vaxart.com.

A replay of the webcast will be available on the Investors page of the Company's website at www.vaxart.com approximately two hours following the conclusion of the event.

As previously announced, Vaxart's 2022 annual meeting of stockholders has been adjourned to Wednesday, July 6, 2022, at 12:30 p.m. Eastern Time. Vaxart encourages all stockholders of record on April 11, 2022, to vote their shares or change their votes in favor of all the proposals being submitted at the annual meeting by 11:59 p.m. Eastern Time on July 5, 2022.

About Vaxart

Vaxart is a clinical-stage biotechnology company developing a range of oral recombinant vaccines based on its proprietary delivery platform. Vaxart vaccines are designed to be administered using tablets that can be stored and shipped without refrigeration and eliminate the risk of needle-stick injury. Vaxart believes that its proprietary tablet vaccine delivery platform is suitable to deliver recombinant vaccines, positioning the company to develop oral versions of currently marketed vaccines and to design recombinant vaccines for new indications. Vaxart's development programs currently include tablet vaccines designed to protect against coronavirus, norovirus, seasonal influenza, and respiratory syncytial virus (RSV), as well as a therapeutic vaccine for human papillomavirus (HPV), Vaxart's first immune-oncology indication. Vaxart has filed broad domestic and international patent applications covering its proprietary technology and creations for oral vaccination using adenovirus and TLR3 agonists.

Note Regarding Forward-Looking Statements

This press release contains forward-looking statements that involve substantial risks and uncertainties. All statements, other than statements of historical facts, included in this press release regarding Vaxart's strategy, prospects, plans and objectives, results from preclinical and clinical trials, commercialization agreements and licenses, and beliefs and expectations of management are forward-looking statements. These forward-looking statements may be accompanied by such words as "should," "believe," "could," "potential," "will," "expected," "anticipate," "plan," and other words and terms of similar meaning. Examples of such statements include, but are not limited to, statements relating to Vaxart's ability to develop and commercialize its product candidates, including its vaccine booster products; Vaxart's expectations regarding clinical results and trial data; and Vaxart's expectations with respect to the effectiveness of its product candidates. Vaxart may not actually achieve the plans, carry out the intentions, or meet the expectations or projections disclosed in the forward-looking statements, and you should not place undue reliance on these forward-looking statements. Actual results or events could differ materially from the plans, intentions, expectations, and projections disclosed in the forward-looking statements. Various important factors could cause actual results or events to differ materially from the forward-looking statements that Vaxart makes, including uncertainties inherent in research and development, including the ability to meet anticipated clinical endpoints, commencement, and/or completion dates for clinical trials, regulatory submission dates, regulatory approval dates, and/or launch dates, as well as the possibility of unfavorable new clinical data and further analyses of existing clinical data; the risk that clinical trial data are subject to differing interpretations and assessments by regulatory authorities; whether regulatory authorities will be satisfied with the design of and results from the clinical studies; decisions by regulatory authorities impacting labeling, manufacturing processes, and safety that could affect the availability or commercial potential of any product candidate, including the possibility that Vaxart's product candidates may not be approved by the FDA or non-U.S. regulatory authorities; that, even if approved by the FDA or non-U.S. regulatory authorities, Vaxart's product candidates may not achieve broad market acceptance; that a Vaxart collaborator may not attain development and commercial milestones; that Vaxart or its partners may experience manufacturing issues and delays due to events within, or outside of, Vaxart's or its partners' control; difficulties in production, particularly in scaling up initial production, including difficulties with production costs and yields, quality control, including stability of the product candidate and quality assurance testing, shortages of qualified personnel or key raw materials, and compliance with strictly enforced federal, state, and foreign regulations; that Vaxart may not be able to obtain, maintain, and enforce necessary patent and other intellectual property protection; that Vaxart's capital resources may be inadequate; Vaxart's ability to resolve pending legal matters; Vaxart's ability to obtain sufficient capital to fund its operations on terms acceptable to Vaxart, if at all; the impact of government healthcare proposals and policies; competitive factors; and other risks described in the "Risk Factors" sections of Vaxart's Quarterly and Annual Reports filed with the SEC. Vaxart does not assume any obligation to update any forward-looking statements, except as required by law.

Contacts

Vaxart Media Relations: Mark Herr Vaxart, Inc. mherr@vaxart.com (203) 517-8957 Investor Relations: Andrew Blazier FINN Partners IR@vaxart.com (646) 871-8486