

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): September 14, 2021

KALVISTA PHARMACEUTICALS, INC.

(Exact name of registrant as specified in its charter)

Delaware  
(State or other jurisdiction  
of incorporation)

001-36830  
(Commission  
File Number)

20-0915291  
(IRS Employer  
Identification No.)

55 Cambridge Parkway  
Suite 901E  
Cambridge, Massachusetts 02142  
(Address of Principal Executive Offices) (Zip Code)

(857) 999-0075  
(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))

Securities registered pursuant to Section 12(b) of the Act:

Title of each class	Trading symbol(s)	Name of each exchange on which registered
Common Stock, \$0.001 Par Value Per Share	KALV	The Nasdaq Stock Market LLC

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 8.01 Other Events.**

On September 14, 2021, KalVista Pharmaceuticals, Inc. (“KalVista”) announced the U.S. Food and Drug Administration (“FDA”) lifted the clinical hold on KalVista’s Phase 2 clinical trial of KVD824 (“KOMLETE”) for oral prophylactic treatment of hereditary angioedema.

The previously announced clinical hold was removed after FDA review of KalVista’s responses to the FDA request for further information and analysis related to certain preclinical studies of KVD824. Refinements were also made to the KVD824 Phase 2 KOMLETE protocol. KalVista is working closely with study investigators and clinical trial sites to proceed with all study activities as soon as possible.

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**SIGNATURE**

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.

Date: September 15, 2021

**KALVISTA PHARMACEUTICALS, INC.**

By: /s/ Benjamin L. Palleiko

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Benjamin L. Palleiko  
Chief Business Officer and Chief Financial Officer