UNITED STATES SECURITIES AND EXCHANGE COMMISSION

Washington, D.C. 20549

FORM 8	-K
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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 8, 2020

ARCUTIS BIOTHERAPEUTICS, INC.

(Exact name of registrant as specified in its charter)

	Delaware (State or other jurisdiction of incorporation)	001-39186 (Commission File Number)	81-2974255 (IRS Employer Identification Number)	
	(Ad	2945 Townsgate Road, Suite 110 Westlake Village, CA 91361 Idress of principal executive offices, including Zip Code	e)	
	Registrant's	telephone number, including area code: (805	5) 418-5006	
	ck the appropriate box below if the Form 8-K filing provisions (see General Instructions A.2. be		ng obligation of the registrant under any of the	
	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))			
Sec	urities registered pursuant to Section 12(b) of the A	Act:		
	Title of each class	Trading Symbol(s)	Name of each exchange on which registered	
C	ommon Stock, par value \$0.0001 per share	ARQT	The Nasdaq Global Select Market	
	cate by check mark whether the registrant is an emoter) or Rule 12b-2 of the Securities Exchange Act		ng 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 8, 2020, Arcutis Biotherapeutics, Inc. (the "Company") issued a press release announcing its plans to advance its program to develop ARQ-151 (topical roflumilast cream) for the treatment of atopic dermatitis into Phase 3 clinical trials following its End-of-Phase 2 meeting with the U.S. Food and Drug Administration, without conducting the previously planned Phase 2b atopic dermatitis trial. The Company anticipates initiating pivotal Phase 3 clinical trials in late 2020 or early 2021.

The foregoing contains "forward-looking" statements, including, among others, statements regarding the expected timing of the initiation of pivotal Phase 3 clinical trials in late 2020 or in early 2021. These statements involve substantial known and unknown risks, uncertainties and other factors that may cause our actual results, levels of activity, performance or achievements to be materially different from the information expressed or implied by these forward-looking statements and you should not place undue reliance on our forward-looking statements. Risks and uncertainties that may cause our actual results to differ include risks inherent in the clinical development process and regulatory approval process, the timing of regulatory filings, and our ability to defend our intellectual property. For a further description of the risks and uncertainties applicable to the Company's business, see the "Risk Factors" section of its Form 10-Q filed with U.S. Securities and Exchange Commission (SEC) on August 11, 2020, as well as any subsequent filings with the SEC. The Company undertake no obligation to revise or update information herein to reflect events or circumstances in the future, even if new information becomes available.

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.

ARCUTIS BIOTHERAPEUTICS, INC.

Date: September 9, 2020

By: /s/ John W. Smither

John W. Smither Chief Financial Officer