# 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) May 11, 2020

# **Allakos Inc.**

(Exact name of registrant as specified in its charter)

Delaware (State or other jurisdiction of incorporation) 001-38582 (Commission File Number) 45-4798831 (IRS Employer Identification No.)

975 Island Drive, Suite 201 Redwood City, California 94065 (Address of principal executive offices, including zip code)

(650) 597-5002

(Registrant's telephone number, including area code)

Not Applicable

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

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, par value \$0.001	ALLK	The Nasdaq Global Select Market

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)

Dere-commencement communications pursuant to Rule 14d-2(b) under the Exchange Act (17 CFR 240.14d-2(b))

Dere-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  $\Box$ 

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 2.02 Results of Operations and Financial Condition.

On May 11, 2020, Allakos Inc. (the "Company") issued a press release reporting its financial results for the first quarter ended March 31, 2020. The full text of the press release is attached as Exhibit 99.1 to this Current Report on Form 8-K and is incorporated herein by reference.

The information in this Item 2.02 of this Form 8-K, including the attached Exhibit 99.1, is intended to be furnished and shall not be deemed "filed" for the purposes of Section 18 of the Securities Exchange Act of 1934, as amended (the "Exchange Act"), or otherwise subject to the liabilities of that section, nor shall it be deemed incorporated by reference in any filing made by the Company under the Securities Act of 1933, as amended, or the Exchange Act, except as expressly set forth by specific reference in such filing.

#### Item 8.01 Other Events.

On May 11. 2020, the Company issued a press release announcing a presentation of antolimab (AK002) in patients with severe allergic conjunctivitis at the 2020 American Society of Cataract and Refractive Surgery Virtual Annual Meeting. The full text of the press release is attached as Exhibit 99.2 to this Current Report on Form 8-K and is incorporated herein by reference.

#### Item 9.01 Financial Statements and Exhibits.

(d) Exhibits

Exhibit Number	Description
99.1	Press Release (First Quarter 2020 Financial Results) dated May 11, 2020.
99.2	Press Release (Announcement of Presentation) dated May 11, 2020.
104	Cover Page Interactive Data File (embedded within the Inline XBRL document)

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## 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.

Allakos Inc.

Date: May 11, 2020

By: /s/ Robert Alexander

Robert Alexander Chief Executive Officer

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### Allakos Reports First Quarter 2020 Financial Results

**REDWOOD CITY, Calif., May 11, 2020** – Allakos Inc. (the "Company") (Nasdaq: ALLK), a biotechnology company developing antolimab (AK002) for the treatment of eosinophil and mast cell-related diseases, today reported financial results for the first quarter ended March 31, 2020.

#### First Quarter 2020 Financial Results

Research and development expenses were \$18.3 million in the first quarter of 2020 as compared to \$15.1 million in the same period in 2019, an increase of \$3.2 million.

General and administrative expenses were \$11.6 million in the first quarter of 2020 as compared to \$5.8 million in the same period in 2019, an increase of \$5.8 million.

Allakos reported a net loss of \$27.8 million in the first quarter of 2020 as compared to \$20.0 million in the same period in 2019, an increase of \$7.8 million. Net loss per basic and diluted share was \$0.57 for the first quarter of 2020 compared to \$0.47 in the same period in 2019.

Allakos ended the first quarter of 2020 with \$479.8 million in cash, cash equivalents and marketable securities.

#### **About Allakos**

Allakos is a late-stage biotechnology company developing antibodies that target immunomodulatory receptors present on immune effector cells involved in allergic, inflammatory, and proliferative diseases. The Company's lead antibody, antolimab (AK002), is being evaluated in a Phase 3 study in eosinophilic gastritis (EG) and/or eosinophilic duodenitis (EoD) and a Phase 2/3 study in eosinophilic esophagitis (EoE). Antolimab targets Siglec-8, an inhibitory receptor selectively expressed on human mast cells and eosinophils. Inappropriately activated eosinophils and mast cells have been identified as key drivers in a number of severe diseases affecting the gastrointestinal tract, eyes, skin, lungs and other organs. Antolimab has been tested in multiple clinical studies, in which antolimab eliminated blood and tissue eosinophils, inhibited mast cells and improved disease symptoms in patients with eosinophilic gastritis and/or eosinophilic duodenitis, eosinophilic esophagitis, mast cell gastrointestinal disease, severe allergic conjunctivitis, chronic urticaria and indolent systemic mastocytosis. For more information, please visit the Company's website at www.allakos.com.

## **Forward-Looking Statements**

This press release contains forward-looking statements within the meaning of The Private Securities Litigation Reform Act of 1995 as contained in Section 27A of the Securities Act of 1933, as amended, and Section 21E of the Securities Exchange Act of 1934, as amended. Such forward-looking statements include, but are not limited to, Allakos' progress and business plans. Such statements are subject to numerous important factors, risks and uncertainties that may cause actual events or results to differ materially from current expectations and beliefs, including but not limited to: Allakos' early stages of clinical drug development; Allakos' ability to timely complete clinical trials for, and if approved, commercialize antolimab (AK002), its lead compound; Allakos' ability to obtain required regulatory approvals for its product candidates; uncertainties related to the enrollment of patients in its clinical trials; Allakos' ability to demonstrate sufficient safety and efficacy of its product candidates in its clinical trials; uncertainties related

to the success of later-stage clinical trials, regardless of the outcomes of preclinical testing and early-stage trials; market acceptance of Allakos' product candidates; uncertainties related to the projections of the size of patient populations suffering from the diseases Allakos is targeting; Allakos' ability to advance additional product candidates beyond antolimab (AK002); Allakos' ability to obtain additional capital to finance its operations; and other important risk factors set forth in Allakos' most recent Annual Report on Form 10-K filed with the SEC on February 25, 2020, Quarterly Report on Form 10-Q filed with the SEC on May 11, 2020 and future reports to be filed with the SEC. These documents contain and identify important factors that could cause the actual results for Allakos to differ materially from those contained in Allakos' forward-looking statements. Any forward-looking statements contained in this press release speak only as of the date hereof, and Allakos specifically disclaims any obligation to update any forward-looking statement, except as required by law. These forward-looking statements should not be relied upon as representing Allakos' views as of any date subsequent to the date of this press release.

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Source: Allakos Inc.

**Investor Contact:** Adam Tomasi, President and COO ir@allakos.com Media Contact: Denise Powell denise@redhousecomms.com

# ALLAKOS INC. STATEMENTS OF OPERATIONS AND COMPREHENSIVE LOSS (in thousands, except per share data) (unaudited)

		Three Months Ended March 31,		
		2020		2019
Operating expenses				
Research and development	\$	18,285	\$	15,098
General and administrative		11,588		5,829
Total operating expenses		29,873		20,927
Loss from operations		(29,873)		(20,927)
Interest income, net		1,989		1,030
Other income (expense), net		60		(56)
Net loss		(27,824)		(19,953)
Unrealized gain on marketable				
securities, net of tax		1,869		45
Comprehensive loss	\$	(25,955)	\$	(19,908)
Net loss per common share:				
Basic and diluted	\$	(0.57)	\$	(0.47)
Weighted-average number of common shares outstanding:				
Basic and diluted		48,691		42,620

# ALLAKOS INC. CONDENSED BALANCE SHEETS (in thousands)

		rch 31, 2020 audited)	D	ecember 31, 2019
Assets	(	,		
Current assets:				
Cash and cash equivalents	\$	135,942	\$	38,367
Investments in marketable securities		343,863		457,534
Prepaid expenses and other current assets		3,953		3,969
Total current assets		483,758		499,870
Property and equipment, net		8,072		8,410
Operating lease right-of-use assets		5,705		5,775
Other long-term assets		2,839		2,839
Total assets	\$	500,374	\$	516,894
Liabilities and stockholders' equity				
Current liabilities:				
Accounts payable	\$	9,062	\$	5,963
Accrued expenses and other current liabilities		5,273		7,098
Total current liabilities		14,335		13,061
Other long-term liabilities		7,995		8,112
Total liabilities		22,330		21,173
Stockholders' equity:				
Common stock		48		48
Additional paid-in capital		693,298		685,020
Accumulated other comprehensive gain		2,006		137
Accumulated deficit		(217,308)		(189,484)
Total stockholders' equity		478,044		495,721
Total liabilities and stockholders' equity	\$	500,374	\$	516,894



#### Allakos Announces a Presentation of Antolimab in Patients with Severe Allergic Conjunctivitis at the 2020 ASCRS Virtual Annual Meeting

REDWOOD CITY, Calif., May 11, 2020 (GLOBE NEWSWIRE) -- Allakos Inc. (the "Company") (Nasdaq: ALLK), a biotechnology company developing antolimab (AK002) for the treatment of eosinophil and mast cell related diseases, today announced the acceptance of an oral presentation at the upcoming American Society of Cataract and Refractive Surgery (ASCRS) Virtual Annual Meeting.

The virtual presentation will take place on May 16, 2020 at 6:00pm ET. The <u>abstract</u> is currently available on the ASCRS website where the virtual presentation will be available as well.

Title (Presenter):	Phase 1b Study of an Anti-Siglec-8 Monoclonal Antibody, Antolimab (AK002), in Severe Allergic Conjunctivitis (Dr.
	Stephen Anesi, MD)
Session (Location):	Corneal Disease (VM - 1, Virtual Room 4)

The Phase 1b clinical trial evaluated antolimab in patients with three forms of severe allergic conjunctivitis refractory to topical treatments; atopic keratoconjunctivitis, vernal keratoconjunctivitis and perennial allergic conjunctivitis. Following the presentation, the slides will be available on the Allakos website.

#### **About Allakos**

Allakos is a late-stage biotechnology company developing antibodies that target immunomodulatory receptors present on immune effector cells involved in allergic, inflammatory, and proliferative diseases. The Company's lead antibody, antolimab (AK002), is being evaluated in a Phase 3 study in eosinophilic gastritis (EG) and/or eosinophilic duodenitis (EoD) and a Phase 2/3 study in eosinophilic esophagitis (EoE). Antolimab targets Siglec-8, an inhibitory receptor selectively expressed on human mast cells and eosinophils. Inappropriately activated eosinophils and mast cells have been identified as key drivers in a number of severe diseases affecting the gastrointestinal tract, eyes, skin, lungs and other organs. Antolimab has been tested in multiple clinical studies, in which antolimab eliminated blood and tissue eosinophils, inhibited mast cells and improved disease symptoms in patients with eosinophilic gastritis and/or eosinophilic duodenitis, eosinophilic esophagitis, mast cell gastrointestinal disease, severe allergic conjunctivitis, chronic urticaria and indolent systemic mastocytosis. For more information, please visit the Company's website at <u>www.allakos.com</u>.

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