# 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) November 2, 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)

 $(650)\ 597\text{-}5002$  (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)			
	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 $\Box$				
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. $\Box$				

#### Item 8.01. Other Events

On November 2, 2020, the Company issued a press release announcing a poster presentation at last week's American College of Gastroenterology 2020 Virtual Annual Scientific Meeting. The press release is filed as Exhibit 99.1 to this Current Report on Form 8-K and is incorporated by reference herein.

#### Item 9.01. Financial Statements and Exhibits.

(d) Exhibits.

Exhibit Number	Description	
99.1	Press Release dated November 2, 2020.	
104	Cover Page Interactive Data File (embedded within the Inline XBRL document)	
	1	

#### 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.

Date: November 2, 2020

Allakos Inc.

By: /s/ Robert Alexander

Robert Alexander, Ph.D.
Chief Executive Officer



## Allakos Announces Presentation from its Eosinophilic Gastrointestinal Diseases Program at the American College of Gastroenterology (ACG) 2020 Annual Scientific Meeting

**REDWOOD CITY, Calif., November 2, 2020** – Allakos Inc. (NASDAQ: ALLK), a biotechnology company developing lirentelimab (AK002) for the treatment of eosinophil and mast cell-related diseases, announced a poster presentation at last week's American College of Gastroenterology 2020 Virtual Annual Scientific Meeting.

The presentation details were as follows:

Title: High Discovery Rate of Previously Undiagnosed Patients with Eosinophilic Gastritis and Duodenitis Using a Systematic

Endoscopic Biopsy Protocol: Screening Data Analysis From ENIGMA, a Randomized Controlled Trial

Presenter: Kathryn A. Peterson, MD, University of Utah School of Medicine

The ePoster (P2797) is available on the ePoster Hall website. In addition, the poster abstract (S1330) appears in the supplement to the October 2020 issue of the American Journal of Gastroenterology.

#### About Eosinophilic Gastritis, Eosinophilic Duodenitis, and Eosinophilic Esophagitis

Eosinophilic gastritis, eosinophilic duodenitis (previously referred to as eosinophilic gastroenteritis), and eosinophilic esophagitis are chronic, often severe, inflammatory diseases characterized by the presence of high levels of eosinophils in the stomach, duodenum, or esophagus, respectively. Common symptoms of the diseases include abdominal pain, nausea, diarrhea, bloating, cramping, early satiety, loss of appetite, vomiting, dysphagia, and weight loss. The current estimated prevalence of eosinophilic gastritis and eosinophilic duodenitis in the United States is approximately 50,000 people. The estimated prevalence of eosinophilic esophagitis in the United States is approximately 150,000 people. The Company believes that these diseases may be significantly under-diagnosed, or misdiagnosed, as other gastrointestinal diseases. There are no treatments approved specifically for these diseases. Treatment with systemic steroids can provide symptomatic improvement, but long-term treatment with steroids is generally not possible due to the numerous side effects. Allakos has received orphan drug designation for lirentelimab in eosinophilic gastritis, eosinophilic gastroenteritis, and eosinophilic esophagitis.

#### **About Allakos**

Allakos is a clinical 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, lirentelimab (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). Lirentelimab 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. Lirentelimab has been tested in multiple clinical studies. In these studies, lirentelimab eliminated blood and tissue eosinophils, inhibited mast cells and improved disease symptoms in patients with EG and/or EoD, EoE, mast cell gastrointestinal disease, severe allergic conjunctivitis, chronic urticaria and indolent systemic mastocytosis. For more information, please visit the Company's website at <a href="https://www.allakos.com">www.allakos.com</a>.

Source: Allakos Inc.

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