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): January 6, 2022

Aligos Therapeutics, Inc.

(Exact name of registrant as specified in its charter)

Delaware (State or other jurisdiction of incorporation)

001-39617 (Commission File Number)

82-4724808 (IRS Employer **Identification Number)**

One Corporate Dr., 2nd Floor South San Francisco, CA 94080 (Address of principal executive offices, including Zip Code)

	Registrant's telepho	ne number, including area code: (800) 466-6059
	appropriate box below if the Form 8-K filing is inte provisions:	ended to simultaneously satisfy the	filing 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))		
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 per share		ALGS	The Nasdaq Stock Market LLC (Nasdaq Global Select Market)
	y check mark whether the registrant is an emerging processor Rule 12b-2 of the Securities Exchange Act of 1934		405 of the Securities Act of 1933 (§230.405 of this
Emerging	growth company $oxtimes$		
	ging growth company, indicate by check mark if the	9	e extended transition period for complying with any e Act. \Box

Item 8.01 Other Events.

On January 6, 2022, Aligos Therapeutics, Inc. (the "Company") announced that it had halted further development of its STOPSTM drug candidate, ALG-010133, in development to address chronic hepatitis B ("CHB"). This decision is based on emerging data from the Phase 1 Study ALG-010133-101 that indicate that at the projected efficacious dose (400 mg, estimated to achieve liver exposures >3 x EC90 for HBsAg inhibition) there is no meaningful HBsAg reduction. Furthermore, higher doses levels (maximum feasible dose is 600 mg) that were planned to be evaluated in a subsequent cohort are very unlikely to reach the 1 log10 IU/mL HBsAg reduction level that the Company had previously defined as necessary to advance the program. No dose limiting safety findings have been identified in CHB subjects dosed at any dose level. Based on this information, Company management reviewed the data with members of the study's Study Review Committee and jointly concluded that these data were not sufficient to support further development of ALG-010133 and that dosing should be discontinued.

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.

ALIGOS THERAPEUTICS, INC.

Date: January 6, 2022 By: <u>/s/ Lesley Ann Calhoun</u>

Lesley Ann Calhoun

Executive Vice President, Chief Financial Officer