UNITED STATES SECURITIES AND EXCHANGE COMMISSION

WASHINGTON, D.C. 20549

		FORM 8-1	K	
		CURRENT REPO	ORT	
Pursuant to Section 13 or 15(d) of the Securities Exchange Act of 1934				
	Date of R	eport (Date of earliest event r	reported): April 3, 2023	
		SSAMER B (Exact name of Registrant as Specific		
Delaware (State or Other Jurisdiction of Incorporation)		001-38796 (Commission File Numb	(II	47-5461709 (IRS Employer Identification No.)
		3013 Science Park R San Diego, California,		
(Address of Principal Executive Offices) (Zip Code) (858) 684-1300 (Registrant's Telephone Number, Including Area Code)				
	k the appropriate box below if the Form 8-K filing wing provisions (see General Instructions A.2. below	-	satisfy the filing obligation of the regis	trant 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))			
Secur	ities registered pursuant to Section 12(b) of the A	ct:		
	Title of each class	Trading Symbol(s)	Name of each exchange on whic	ch registered
	Common Stock, \$0.0001 par value per share	GOSS	Nasdaq Global Select M	Market
	ate by check mark whether the registrant is an emo er) or Rule 12b-2 of the Securities Exchange Act			f 1933 (§230.405 of this
Emer	ging growth company □			
	emerging growth company, indicate by check mar rised financial accounting standards provided purs			for complying with any new

Item 8.01 Other Events.

On April 3, 2023, Gossamer Bio, Inc. (the "Company") announced that the U.S. Food and Drug Administration (the "FDA") has placed a partial clinical hold on all trials of GB5121, in response to serious adverse events that have been observed in the Phase 1b/2 STAR-CNS study, including atrial fibrillation, a sudden death event and a fatal intracranial hemorrhage. The Company previously announced on March 17, 2023 that it had paused enrollment of the Phase 1b/2 STAR-CNS Study based on the benefit / risk profile observed to date and a prioritization of resources to support the Company's seralutinib program. Based on the foregoing factors, the Company has decided to terminate all ongoing studies and discontinue development of GB5121.

SIGNATURES

Pursuant to the requirements of the Securities Exchange Act of 1934, the registrant has duly caused this report to be signed on its behalf by the undersigned thereunto duly authorized.

Date: April 3, 2023

GOSSAMER BIO, INC.

By: /s/ Christian Waage

Christian Waage

Executive Vice President, Technical Operations & Administration