UNITED STATES SECURITIES AND EXCHANGE COMMISSION WASHINGTON, DC 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): July 8, 2021 (July 8, 2021)

PLURISTEM THERAPEUTICS INC.

(Exact Name of Registrant as Specified in Its Charter)

Nevada	001-31392	98-0351734
(State or Other Jurisdiction of Incorporation)	(Commission File Number)	(IRS Employer Identification No.)
-		racinimental (vo.)
MATAM Advanced Technology	Park	
Building No. 5 Haifa, Israel		3508409
(Address of Principal Executive Of	fices)	(Zip Code)
	011 972 74 710 7171 (Registrant's telephone number, including area code)	
	Not applicable	
(Fo	rmer name or former address, if changed since last repo	ort)
Check the appropriate box below if the Form 8-K filiprovisions:	ing is intended to simultaneously satisfy the filing of	bligation of the registrant under any of the following
☐ Written communications pursuant to Rule 425 under the Securities Act (17 CFR 230.425)		
☐ Soliciting material pursuant to Rule 14a-12 under th	e Exchange Act (17 CFR 240.14a-12)	
☐ Pre-commencement communications pursuant to Ru	ale 14d-2(b) under the Exchange Act (17 CFR 240.14d-	2(b))
☐ Pre-commencement communications pursuant to Ru	ale 13e-4(c) under the Exchange Act (17 CFR 240.13e-4	ł(c))
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 \$0.00001 per share	PSTI	The Nasdaq Global Market
Indicate by check mark whether the registrant is an em chapter) or Rule 12b-2 of the Securities Exchange Act of Emerging growth company □ If an emerging growth company, indicate by check mar	f 1934 (§240.12b-2 of this chapter). k if the registrant has elected not to use the extended to	
financial accounting standards provided pursuant to Sect	ion 13(a) of the Exchange Act. ⊔	

Item 8.01. Other Events.

On July 8, 2021, Pluristem Therapeutics Inc., or the registrant, announced that it is bringing its Acute Respiratory Distress Syndrome, or ARDS associated with COVID-19 Phase II studies to clinical readout. The analysis will be based on 89 patients enrolled in the previously announced two Phase II studies, which investigated the safety and efficacy of the registrant's PLX cells as a treatment for severe COVID-19 cases complicated by ARDS. The registrant expects to announce the topline results of the readout during the fourth quarter of 2021. The registrant also announced that it will not pursue plans announced in December 2020 to expand its COVID-19 program to Mexico in collaboration with Innovare R&D.

Warning Concerning Forward Looking Statements

This Current Report on Form 8-K contains forward-looking statements within the meaning of the "safe harbor" provisions of the Private Securities Litigation Reform Act of 1995 and federal securities laws. For example, forward-looking statements are being used when the registrant discusses the timing of the Phase II study clinical readout. These forward-looking statements and their implications are based on the current expectations of the management of the registrant only, and are subject to a number of factors and uncertainties that could cause actual results to differ materially from those described in the forward-looking statements. The following factors, among others, could cause actual results to differ materially from those described in the forward-looking statements: changes in technology and market requirements; the registrant may encounter delays or obstacles in launching and/or successfully completing its clinical trials; the registrant's products may not be approved by regulatory agencies, the registrant's technology may not be validated as it progresses further and its methods may not be accepted by the scientific community; the registrant may be unable to retain or attract key employees whose knowledge is essential to the development of our products; unforeseen scientific difficulties may develop with our process; the registrant's products may wind up being more expensive than the registrant anticipates; results in the laboratory may not translate to equally good results in real surgical settings; results of preclinical studies may not correlate with the results of human clinical trials; the registrant's patents may not be sufficient; the registrant's products may harm recipients; changes in legislation; inability to timely develop and introduce new technologies, products and applications; loss of market share and pressure on pricing resulting from competition, which could cause the actual results or performance of the registrant to differ materially from those contemplated in such forward-looking statements. In addition, historic results of scientific research do not guarantee that the conclusions of future research would not suggest different conclusions or that historic results would not be interpreted differently in light of additional research or otherwise. Except as otherwise required by law, the registrant undertakes no obligation to publicly release any revisions to these forward-looking statements to reflect events or circumstances after the date hereof or to reflect the occurrence of unanticipated events. For a more detailed description of the risks and uncertainties affecting the registrant, reference is made to the registrant's reports filed from time to time with the Securities and Exchange Commission.

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 hereunto duly authorized.

PLURISTEM THERAPEUTICS INC.

Date: July 8, 2021 By: /s/ Chen Franco-Yehuda

Name: Chen Franco-Yehuda Title: Chief Financial Officer