UNITED STATES SECURITIES AND EXCHANGE COMMISSION

Washington, D.C. 20549

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

Poseida Therapeutics, Inc. (Exact name of registrant as specified in its charter)

Delaware (State or other jurisdiction of incorporation)		001-39376 (Commission File Number)	47-2846548 (I.R.S. Employer Identification No.)
	9390 Towne Centre Drive, S	Suite 200	
San Diego, Californ			92121
	(Address of principal executive	offices)	(Zip Code)
	Registrant's telepho	one number, including area code: (85	58) 779-3100
	(Former nam	\mathbf{N}/\mathbf{A} ne or former address, if changed since last rep	ort.)
	appropriate box below if the Form 8-K filing is int provisions:	ended to simultaneously satisfy the fil	ing 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(s)	Name of each exchange on which registered
Common stock, par value \$0.0001 per share		PSTX	Nasdaq Global Select Market
	y check mark whether the registrant is an emerging f this chapter) or Rule 12b–2 of the Securities Exch		
Emerging	growth company ⊠		
	ging growth company, indicate by check mark if th	•	1 100

Item 8.01 Other Events.

On November 2, 2020, the Company announced that the U.S. Food and Drug Administration (FDA) has lifted a clinical hold on the Company's Phase 1 study of P-PSMA-101 in metastatic castration-resistant prostate cancer (mCRPC) and the Company plans to resume the trial immediately. P-PSMA-101 is the Company's first solid tumor autologous CAR-T therapeutic candidate.

The Company has agreed to implement protocol amendments intended to increase patient compliance and safety that include modified inclusion and exclusion criteria and frequency of monitoring and laboratory testing.

Forward-Looking Statements

Statements contained in this report regarding matters that are not historical facts are "forward-looking statements" within the meaning of the Private Securities Litigation Reform Act of 1995. Because such statements are subject to risks and uncertainties, actual results may differ materially from those expressed or implied by such forward-looking statements. These forward-looking statements are based upon the Company's current expectations and involve assumptions that may never materialize or may prove to be incorrect. Actual results could differ materially from those anticipated in such forward-looking statements as a result of various risks and uncertainties, which include, without limitation, risks and uncertainties associated with development and regulatory approval of novel product candidates in the biopharmaceutical industry and the other risks described in the Company's filings with the Securities and Exchange Commission. All forward-looking statements contained in this report speak only as of the date on which they were made. The Company undertakes no obligation to update such statements to reflect events that occur or circumstances that exist after the date on which they were made, except as required by law.

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.

Poseida Therapeutics, Inc.

Date: November 2, 2020 By: /s/ Harry J. Leonhardt

Harry J. Leonhardt

General Counsel and Chief Compliance Officer