UNITED STATES SECURITIES AND EXCHANGE COMMISSION

Washington, D.C. 20549

		FORM 8-K			
		Current Report or 15(d)of the Securitie Date of earliest event reporte	ies Exchange Act of 1934 ted): May 16, 2022		
		ERE THERAPEUTIC ne of registrant as specified i	•		
	Delaware (State or other jurisdiction of incorporation)	001-36257 (Commission File Number)	27-4842691 (I.R.S. Employer Identification No.)		
		11 Valley Centre Drive, Suite San Diego, CA 92130 incipal Executive Offices, inc			
	(Registrant's	(888) 969-7879 s Telephone Number, includir	ing Area Code)		
	(Former Name or	Not Applicable Former Address, if Changed	d Since Last Report)		
	ck the appropriate box below if the Form 8-K filing is inving provisions:	rended to simultaneously sati	tisfy 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))				
Secu	urities 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	TVTX	The Nasdaq Global Market		
chap	ter) or Rule 12b-2 of the Securities Exchange Act of 19		ned in Rule 405 of the Securities Act of 1933 (§230.405 of the oter).	his	
f an	rging growth company \square emerging growth company, indicate by check mark if t or revised financial accounting standards provided pur		t to use the extended transition period for complying with a exchange Act. \Box	ny	

Item 8.01 Other Events.

On May 16, 2022, Travere Therapeutics, Inc. (the "Company") announced that the U.S. Food and Drug Administration ("FDA") has accepted and granted Priority Review of its New Drug Application ("NDA") under Subpart H for accelerated approval of sparsentan for the treatment of IgA nephropathy ("IgAN"). The FDA has indicated that it is not currently planning to hold an advisory committee meeting to discuss the application and has assigned a Prescription Drug User Fee Act (PDUFA) target action date of November 17, 2022.

Forward-Looking Statements

This Current Report on Form 8-K contains "forward-looking statements" as that term is defined in the Private Securities Litigation Reform Act of 1995. Such forward-looking statements include references relating to the FDA's potential approval of the Company's NDA for sparsentan for the treatment of IgAN by the target action date of November 17, 2022, or at all, and the expectation around any potential future request by the FDA to hold an advisory committee meeting related to the application. Such forward-looking statements are based on current expectations and involve inherent risks and uncertainties, including factors that could delay, divert or change any of them, and could cause actual outcomes and results to differ materially from current expectations. No forward-looking statement can be guaranteed. Among the factors that could cause actual results to differ materially from those indicated in the forward-looking statements are risks and uncertainties associated with the regulatory review and approval process, including the Subpart H accelerated approval pathway. There is no guarantee that the FDA will grant accelerated approval of sparsentan for IgAN or that sparsentan will be approved at all. In addition, such risks and uncertainties may include those described in the Company's filings with the Securities and Exchange Commission, including under the "Risk Factors" heading of the Company's quarterly report on Form 10-Q for the quarter ended March 31, 2022, as filed with the SEC on May 5, 2022. You are cautioned not to place undue reliance on any forward-looking statements as there are important factors that could cause actual results to differ materially from those in any forward-looking statements, many of which are beyond our control. Except to the extent required by law, the Company undertakes no obligation to publicly update any forward-looking statement.

Item 9.01 Financial Statements and Exhibits.

(d) Exhibits

Exhibit No.	Description
104	Cover Page Interactive Data File (embedded within the Inline XBRL document).

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.

TRAVERE THERAPEUTICS, INC.

By: /s/ Elizabeth E. Reed

Dated: May 16, 2022

Name: Elizabeth E. Reed

Title: Senior Vice President, General Counsel and Secretary