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): November 11, 2020

Lineage Cell Therapeutics, Inc. (Exact name of registrant as specified in charter)

001-12830

(Commission

File Number)

94-3127919

(IRS Employer

Identification No.)

California

(State or other jurisdiction

of incorporation)

2173 Salk Avenue, Sui Carlsbad, Californ (Address of principal executi	ia	92008 (Zip Code)
	(442) 287-8990 Registrant's telephone number, inclu	uding area code
	(Former name or former address, if chang	ged since last report)
Check the appropriate box below if the Form following provisions (see General Instruction A		sly satisfy the filing obligation of the registrant under any of the
☐ Written communications pursuant to Rule	425 under the Securities Act (17 CFR 230	0.425)
☐ Soliciting material pursuant to Rule 14a-12	under the Exchange Act (17 CFR 240.1	4a-12)
☐ Pre-commencement communications pursu	uant to Rule 14d-2(b) under the Exchange	Act (17 CFR 240.14d-2(b))
☐ Pre-commencement communications pursu	uant to Rule 13e-4(c) under the Exchange	Act (17 CFR 240.13e-4(c))
	Securities registered pursuant to Section	n 12(b) of the Act:
Title of each class	Trading Symbol(s)	Name of each exchange on which registered
Common stock	LCTX	NYSE American
Indicate by check mark whether the registrant i of this chapter) or Rule 12b-2 of the Securities		ed in as defined in Rule 405 of the Securities Act of 1933 (§230.405 is chapter).
		Emerging growth company \Box
If an emerging growth company, indicate by clor revised financial accounting standards provide		at to use the extended transition period for complying with any new lange Act. \Box
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Item 8.01. Other Events

On November 11, 2020, Lineage Cell Therapeutics, Inc. announced that it completed enrollment in its Phase 1/2a multicenter clinical trial of OpRegen[®], a retinal pigment epithelium cell transplant therapy for the treatment of advanced dry age-related macular degeneration with geographic atrophy. The trial enrolled a total of 24 patient into four cohorts. Seven of the patients were dosed with the 510(k)-cleared Subretinal Delivery System developed by Gyroscope Therapeutics.

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.

Lineage Cell Therapeutics, Inc.

Date: November 12, 2020 By: /s/ Chase C. Leavitt

Name: Chase C. Leavitt

Title: General Counsel and Corporate Secretary