

**UNITED STATES  
SECURITIES AND EXCHANGE COMMISSION**  
Washington, D.C. 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 4, 2020**

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

**Delaware**  
(State or other jurisdiction  
of incorporation)

**001-39059**  
(Commission  
File Number)

**85-1021707**  
(IRS Employer  
Identification No.)

**28159 Avenue Stanford, Suite 220, Valencia, CA 91355**  
(Address of principal executive offices, including Zip Code)

**661.367.9170**  
(Registrant's telephone number, including area code)

**N/A**  
(Former name or former address, if changed since last report)

Check the appropriate box below if the Form 8-K filing is intended to simultaneously satisfy the filing obligation of the registrant under any of the following provisions:

- 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	RCEL	The Nasdaq Stock Market LLC

Indicate by check mark whether the registrant is an emerging growth company as defined in Rule 405 of the Securities Act of 1933 or Rule 12b-2 of the Securities Exchange Act of 1934. Emerging growth company

If an emerging growth company, indicate by check mark if the registrant has elected not to use the extended transition period for complying with any new or revised financial accounting standards provided pursuant to Section 13(a) of the Exchange Act.

If an emerging growth company, indicate by check mark if the registrant has elected not to use the extended transition period for complying with any new or revised financial accounting standards provided pursuant to Section 13(a) of the Exchange Act.

### **Item 8.01. Other Events**

On November 4, 2020, AVITA Therapeutics, Inc. (the Company) issued a press release announcing a correction in the time for the adjourned 2020 Annual Stockholder Meeting (the Adjourned Annual Meeting), due to the change in daylight savings time Pacific Time in the United States, it results in change in time in Australian Eastern Daylight Time (AEDT) from 7:00 am to 8:00 am on Tuesday, November 10, 2020 and remains unchanged at 1:00 pm Pacific Time on Monday, November 9, 2020.

Therefore, the Adjourned Annual Meeting will be held 8:00 am (AEDT) on Tuesday, November 10, 2020 and 1:00 pm (Pacific Time) on Monday, November 9, 2020.

Stockholders who have any questions or require any assistance with how to complete a proxy or CDI Voting Instruction Form or who do not have the required materials, may contact Okapi Partners by telephone (toll-free within North America) at (877) 629-6356 or (call collect outside North America) at (212) 297-0720 or by email at [info@okapipartners.com](mailto:info@okapipartners.com).

### **Additional Information and Where to Find It**

This document may be deemed to be solicitation material in respect of the Annual Meeting to be held on November 9, 2020 (Pacific Time) (being November 10, 2020 (AEDT)). In connection with the Annual Meeting, the Company filed a definitive proxy statement with the SEC on September 25, 2020. BEFORE MAKING ANY VOTING DECISIONS, STOCKHOLDERS AND CDI HOLDERS ARE URGED TO READ THE DEFINITIVE PROXY STATEMENT AND ANY OTHER RELEVANT DOCUMENTS FILED WITH THE SEC OR THE AUSTRALIAN SECURITIES EXCHANGE (“ASX”), BECAUSE THEY CONTAIN IMPORTANT INFORMATION ABOUT THE ANNUAL MEETING. The definitive proxy statement was mailed to stockholders and CDI holders entitled to vote at the Annual Meeting. No changes have been made to the proposals to be voted on by stockholders at the Annual Meeting. The Company’s proxy statement and any other materials filed by the Company with the SEC can be obtained free of charge at the SEC’s website at <https://www.sec.gov/Archives/edgar/data/1762303/000119312520254171/d37466ddef14a.htm> or on the Company’s website at [www.avitamedical.com](http://www.avitamedical.com). The proxy statement and any other materials filed by the Company with the ASX can be obtained free of charge at the ASX’s website [www.asx.com.au](http://www.asx.com.au) and also the Company’s website at [www.avitamedical.com](http://www.avitamedical.com).

### **Item 9.01. Financial Statements and Exhibits.**

(d) Exhibits.

#### **Exhibit No.    Description**

99.1    [Press release of AVITA Therapeutics, Inc. dated November 4, 2020](#)

**SIGNATURES**

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

**AVITA THERAPEUTICS, INC.**  
(Registrant)

Date: November 4, 2020

*/s/ David McIntyre*

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Name: David McIntyre

Title: Chief Financial Officer



**AVITA THERAPEUTICS, INC. ANNOUNCES CORRECTION OF TIME FOR THE  
ADJOURNED  
2020 ANNUAL STOCKHOLDER MEETING**

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**Valencia, Calif, USA, November 4, 2020/Melbourne, November 5, 2020** — AVITA Therapeutics, Inc. (NASDAQ: RCEL, ASX:AVH), a regenerative medicine company that is developing and commercializing a technology platform that enables point-of-care autologous skin restoration for multiple unmet needs, announces today a correction in the time for the adjourned 2020 Annual Stockholder Meeting (“Adjourned Annual Meeting”), due to the change in daylight savings time Pacific Time in the United States, it results in change in time in Australian Eastern Daylight Time (AEDT) from 7:00 am to 8:00 am on Tuesday, November 10, 2020 and remains unchanged at 1:00 pm Pacific Time on Monday, November 9, 2020.

Therefore, the Adjourned Annual Meeting will be held 8:00 am (AEDT) on Tuesday, November 10, 2020 and 1:00 pm (Pacific Time) on Monday, November 9, 2020.

Stockholders who have any questions or require any assistance with how to complete a proxy or CDI Voting Instruction Form or who do not have the required materials, may contact Okapi Partners by telephone (toll-free within North America) at (877) 629-6356 or (call collect outside North America) at (212) 297-0720 or by email at [info@okapipartners.com](mailto:info@okapipartners.com).

Authorized for release by the Chief Executive Officer of AVITA Therapeutics, Inc.

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**Important Information**

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AVITA Therapeutics, Inc. (ARBN 641 28 155) c/o Mertons Corporate Services Pty Ltd Level 7, 330 Collins Street, Melbourne Victoria 3000

## ABOUT AVITA THERAPEUTICS, INC.

AVITA Therapeutics is a regenerative medicine company with a technology platform positioned to address unmet medical needs in burns, chronic wounds, and aesthetics indications. AVITA Therapeutics' patented and proprietary collection and application technology provides innovative treatment solutions derived from the regenerative properties of a patient's own skin. The medical devices work by preparing a RES® REGENERATIVE EPIDERMAL SUSPENSION, an autologous suspension comprised of the patient's skin cells necessary to regenerate natural healthy epidermis. This autologous suspension is then sprayed onto the areas of the patient requiring treatment.

AVITA Therapeutics' first U.S. product, the RECELL® System, was approved by the U.S. Food and Drug Administration (FDA) in September 2018. The RECELL System is indicated for use in the treatment of acute thermal burns in patients 18 years and older. The RECELL System is used to prepare Spray-On Skin™ Cells using a small amount of a patient's own skin, providing a new way to treat severe burns, while significantly reducing the amount of donor skin required. The RECELL System is designed to be used at the point of care alone or in combination with autografts depending on the depth of the burn injury. Compelling data from randomized, controlled clinical trials conducted at major U.S. burn centers and real-world use in more than 8,000 patients globally, reinforce that the RECELL System is a significant advancement over the current standard of care for burn patients and offers benefits in clinical outcomes and cost savings. Healthcare professionals should read the INSTRUCTIONS FOR USE—RECELL® Autologous Cell Harvesting Device (<https://recellsystem.com/>) for a full description of indications for use and important safety information including contraindications, warnings and precautions.

In international markets, our products are marketed under the RECELL System brand to promote skin healing in a wide range of applications including burns, chronic wounds and aesthetics. The RECELL System is TGA-registered in Australia and received CE-mark approval in Europe. To learn more, visit [www.avitamedical.com](http://www.avitamedical.com).

## CAUTIONARY NOTE REGARDING FORWARD-LOOKING STATEMENTS

*This letter includes forward-looking statements. These forward-looking statements generally can be identified by the use of words such as “anticipate,” “expect,” “intend,” “could,” “may,” “will,” “believe,” “estimate,” “look forward,” “forecast,” “goal,” “target,” “project,” “continue,” “outlook,” “guidance,” “future,” other words of similar meaning and the use of future dates. Forward-looking statements in this letter include, but are not limited to, statements concerning, among other things, our ongoing clinical trials and product development activities, regulatory approval of our products, the potential for future growth in our business, and our ability to achieve our key strategic, operational and financial goal. Forward-looking statements by their nature address matters that are, to different degrees, uncertain. Each forward-looking statement contained in this letter is subject to risks and uncertainties that could cause actual results to differ materially from those expressed or implied by such statement. Applicable risks and uncertainties include, among others, the timing of regulatory approvals of our products; physician acceptance, endorsement, and use of our products; failure to achieve the anticipated benefits from approval of our products; the effect of regulatory actions; product liability claims; risks associated with international operations and expansion; and other business effects, including the effects of industry, economic or political conditions outside of the company's control. Investors should not place considerable reliance on the forward-looking statements contained in this letter. Investors are encouraged to read our publicly available filings for a discussion of these and other risks and uncertainties. The forward-looking statements in this letter speak only as of the date of this release, and we undertake no obligation to update or revise any of these statements.*

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**FOR FURTHER INFORMATION:**

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