ASX/News Release

AVITA MEDICAL, INC. (ASX:AVH)

Cleansing Notice under section 708A(5)(e) of the Corporations Act 2001 (Cth)

Valencia, Calif., USA, 15 February 2024 and Melbourne, Australia, 16 February 2024: On 9 February 2024 (United States) / 10 February 2024 (Australia), AVITA Medical, Inc. (**Company**) issued a total of 17,963 fully paid shares of common stock in the Company (**New Securities**). 2,234 shares were issued as a result of the exercise of 2,234 unquoted Options. 15,729 shares were issued as a result of the exercise of 1,572,900 unquoted Options in AVITA Medical Pty Limited (Avita Australia) (which entitled the shareholder to be issued shares of common stock in the Company rather than ordinary shares in Avita Australia on a consolidation ratio of 100:1 as set out in the Company's pre-quotation disclosure released to the market on 24 June 2020).

The New Securities will be quoted on NASDAQ, but may be converted into CHESS Depositary Interests (**CDIs**) in the Company quoted on ASX at any time by the relevant holder. The Company seeks to rely on an exemption under section 708A of the *Corporations Act 2001* (Cth) (**Corporations Act**) with respect to the sale of any CDIs which are issued on conversion of the New Securities (in the instance that such conversion occurs).

The Company gives this notice under section 708A(5)(e) of the Corporations Act as modified by ASIC Class Order 14/827.

The New Securities were issued without disclosure to investors under Part 6D.2 of the Corporations Act.

As at the date of this notice, the Company has complied with:

- the provisions of Chapter 2M of the Corporations Act as they apply to the Company; and
- section 674 of the Corporations Act.

As at the date of this notice, there is no information that is 'excluded information' within the meaning of section 708A(7) and section 708A(8) of the Corporations Act.

Authorized for release by the Chief Financial Officer of AVITA Medical, Inc.

ABOUT AVITA MEDICAL, INC.

AVITA Medical® is a regenerative medicine company leading the development and commercialization of devices and autologous cellular therapies for skin restoration. The RECELL® System technology platform, approved by the FDA for the treatment of thermal burn wounds and full-thickness skin defects and for repigmentation of stable depigmented vitiligo lesions, harnesses the regenerative properties of a patient's own skin to create Spray-On Skin™ cells. Delivered at the point-of-care, RECELL enables improved clinical outcomes. RECELL is the catalyst of a new treatment paradigm and AVITA Medical is leveraging its proven and differentiated capabilities to develop first-in-class cellular therapies for multiple indications.

In international markets, our products are approved under the RECELL System brand to promote skin healing in a wide range of applications including burns, soft tissue repair, vitiligo, and aesthetics. The RECELL System is TGA-registered in Australia, received CE-mark approval in Europe and has PMDA approval in Japan.

AVITA Medical, Inc. Suite 220, 28159 Avenue Stanford, Valencia, California 91355 United States of America

CAUTIONARY NOTE REGARDING FORWARD-LOOKING STATEMENTS

This announcement 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 announcement 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 goals. Forward-looking statements by their nature address matters that are, to different degrees, uncertain. Each forward- looking statement contained in this announcement 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 announcement. 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 announcement speak only as of the date of this announcement, and we undertake no obligation to update or revise any of these statements.

FOR FURTHER INFORMATION:

Investors & Media
AVITA Medical, Inc.
Jessica Ekeberg
Phone +1-661-904-9269
investor@avitamedical.com
media@avitamedical.com

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