



AVITA MEDICAL, INC. (ASX:AVH)

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

Valencia, Calif., USA, 13 November 2024 and Melbourne, Australia, 14 November 2024: On 8 November 2024 (United States) / 9 November 2024 (Australia) and 12 November 2024 (United States) / 13 November 2024 (Australia) AVITA Medical, Inc. (**Company**) issued a total of 12,117 fully paid shares of common stock in the Company (**New Securities**). 9,217 securities were issued as a result of the exercise of 9,217 unquoted Options. 2,900 securities were issued as a result of the exercise of 290,000 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-quotations disclosure released to the market on 24 June 2020).

The New Securities will be quoted on NASDAQ, but may be converted into CHESSE 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 and 674A 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 commercial-stage regenerative medicine company transforming the standard of care in wound care management and skin restoration with innovative devices. At the forefront of our platform is the RECELL® System, approved by the Food and Drug Administration for the treatment of thermal burn wounds and full-thickness skin defects, and for repigmentation of stable depigmented vitiligo lesions. RECELL harnesses the regenerative properties of a patient's own skin to create Spray-On Skin™ Cells, delivering a transformative solution at the point-of-care. This breakthrough technology serves as the catalyst for a new treatment paradigm enabling improved clinical outcomes. AVITA Medical also holds the exclusive rights to market, sell, and distribute PermeaDerm®, a biosynthetic wound matrix, in the United States.

In international markets, the RECELL System is approved to promote skin healing in a wide range of applications including burns, full-thickness skin defects, and vitiligo. The RECELL System is TGA-registered in Australia, received CE-mark approval in Europe and has PMDA approval in Japan.

To learn more, visit www.avitamedical.com.

Forward-Looking Statements

Statements in this press release may contain forward-looking statements within the meaning of the Private Securities Litigation Reform Act of 1995. Such forward-looking statements are subject to significant risks and uncertainties that could cause actual results to differ materially from those expressed or implied by such statements. Forward-looking statements generally may 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,” and similar words or expressions, and the use of future dates. Applicable risks and uncertainties include, among others, the timing and realization 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. These statements are made as of the date of this release, and the Company undertakes no obligation to publicly update or revise any of these statements, except as required by law. For additional information and other important factors that may cause actual results to differ materially from forward-looking statements, please see the “Risk Factors” section of the Company’s latest Annual Report on Form 10-K and other publicly available filings for a discussion of these and other risks and uncertainties.

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