RECELL® System Data to be Presented at the Controversies and Conversations in Laser & Cosmetic Surgery Annual Meeting

August 17, 2022

Two presentations will highlight data utilizing the RECELL platform for repigmentation

VALENCIA, Calif. and MELBOURNE, Australia, Aug. 17, 2022 (GLOBE NEWSWIRE) -- AVITA Medical, 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, announced today that a poster presentation on cell characterization and potential clinical benefits of the RECELL® Autologous Cell Suspension System (RECELL® System) for the treatment of stable vitiligo will be shared at the Controversies and Conversations in Laser & Cosmetic Surgery Annual Meeting. The conference will be held in Santa Barbara, CA, on August 19-21 bringing experts together to discuss controversial issues in cutaneous and aesthetic surgery or challenging therapeutic problems within dermatology.

“Given the unique format of this meeting, we look forward to the presentation of RECELL and, more importantly, the conversation amongst dermatology experts as they discuss the unique attributes of this platform, including the potential for in-office point-of-care treatment in about 30 minutes,” said Dr. Mike Perry, Chief Executive Officer of AVITA Medical. “Following review by the FDA, we believe the RECELL System may well offer a welcome treatment option for patients seeking repigmentation for stable vitiligo lesions.”

RECELL® System Presentations

- Autologous Melanocyte Transfer for Repigmentation and Scar Healing – Girish Munavalli, MD, MHS
- **Characterization of Autologous Skin Cell Suspension prepared using RECELL® Autologous Cell Suspension System and Evidence of Melanocyte Transfer** – G. Kashgari, et al

In the U.S., the RECELL® System is indicated for the treatment of acute thermal burn wounds in patients 18 years of age and older or application in combination with meshed autografting for acute full-thickness thermal burn wounds in pediatric and adult patients. Physician-initiated research beyond the FDA approved indication is not sponsored by AVITA Medical and contains independent data.

AVITA Medical is currently completing a pivotal trial for the use of the RECELL System for treatment of stable vitiligo. The vitiligo clinical trial aims to demonstrate safe and effective repigmentation when using the RECELL System in combination with phototherapy. AVITA anticipates FDA approval in 2023.

ABOUT AVITA MEDICAL, INC.

AVITA Medical, Inc. is a regenerative medicine company with a technology platform positioned to address unmet medical needs in burns, chronic wounds, and aesthetics indications. AVITA Medical Inc.’s patented, and proprietary collection and application technology provides innovative treatment solutions derived from the regenerative properties of a patient’s own skin. The Company’s lead product is the RECELL® System, a device that enables healthcare professionals to Spray-On Skin™ Cells using a small sample of the patient’s own skin to create an autologous suspension. The RES® Regenerative Epidermal Suspension™ is then sprayed onto the areas of the patient requiring treatment to regenerate natural healthy epidermis.

AVITA Medicals’ 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 approved for acute partial-thickness thermal burn wounds in patients 18 years of age and older or application in combination with meshed autografting for acute full-thickness thermal burn wounds in pediatric and adult patients. In February 2022, the FDA reviewed and approved the PMA supplement for RECELL Autologous Cell Harvesting Device, an enhanced RECELL System aimed at providing clinicians a more efficient user experience and simplified workflow.

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 15,000 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 approved 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, received CE-mark approval in Europe, and received Japan’s Pharmaceuticals and Medical Devices Act (PMDA) approval for burns in Japan.

CAUTIONARY NOTE REGARDING FORWARD-LOOKING STATEMENTS

This press release 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 press release 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 press release 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 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. Investors should not place considerable reliance on the forward-looking statements contained in this press release. 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 press release speak only as of the date of this release, and we undertake no obligation to update or revise any of these statements.

This press release was authorized by the review committee of AVITA Medical, Inc.

FOR FURTHER INFORMATION:

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