ASX/News Release

AVITA Medical Announces RECELL® System Preliminary Data Focused on Vitiligo and Facial Acne Scars at the 24th Annual World Congress of Dermatology

Abstracts Showcase Research on Potential Use of RECELL® System in Regenerative Dermatology

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Valencia, Calif., USA, and Melbourne, Australia, June 12, 2019 —AVITA Medical (ASX: AVH, OTCQX: AVMXY), a regenerative medicine company with a technology platform positioned to address unmet medical needs in therapeutic skin restoration, announces new preliminary RECELL® Autologous Cell Harvesting Device (RECELL® System) data at the 24th annual World Congress of Dermatology Meeting June 10-15, 2019, in Milan, Italy. Three posters at the World Congress of Dermatology will highlight the potential use of the RECELL® System for the treatment of vitiligo and facial acne scars.

"Building on the foundation of long-term safety and efficacy of the RECELL® System for the treatment of burn patients, we look forward to presenting data at the World Congress of Dermatology exploring the use of this innovative therapy for regenerative dermatology," said Dr. Michael Perry, AVITA Medical's Chief Executive Officer. "We are committed to pursuing the full potential of the RECELL® System as a meaningful treatment option to advance patient care in areas with significant unmet medical need."

The RECELL® System, which uses a small amount of a patient's own skin to prepare Spray-On Skin™ Cells at the point of care, is approved by the U.S. Food and Drug Administration (FDA) for the treatment of acute thermal burns in patients 18 years and older. In international markets, the RECELL® System is approved to promote skin healing in a wide range of applications including burns, acute wounds, scars and vitiligo.

RECELL® System Poster Presentations at World Congress of Dermatology

Poster Exhibition Date, Time (CET), Location	**Poster Title**	**Authors**
June 11, 7:00 a.m. – 6:30 p.m. June 12, 7:00 a.m. – 7:30 p.m. June 13, 7:00 a.m. – 7:30 p.m. June 14, 7:00 a.m. – 7:00 p.m. June 15, 7:00 a.m. – 1:00 p.m. MiCo – Milano Congressi South Wing, Level 0	"Treatment of Facial Acne Scars in Chinese Patients: Combination of Dermabrasion and ReCell Technique."	**Nanze Yu, et al** Peking Union Medical College Hospital Beijing, China
June 11, 7:00 a.m. – 6:30 p.m. June 12, 7:00 a.m. – 7:30 p.m. June 13, 7:00 a.m. – 7:30 p.m. June 14, 7:00 a.m. – 7:00 p.m.	"Is Suction Blister Epidermal Grafting a Simple and Reliable Way to Screen Patients with Large Area Vitiligo for ReCell Treatment?"	**Weiwei Li, et al** Beijing Tsinghua Changgung Hospital Beijing, China

June 15, 7:00 a.m. – 1:00 p.m. MiCo – Milano Congressi South Wing, Level 0		
June 11, 7:00 a.m. – 6:30 p.m. June 12, 7:00 a.m. – 7:30 p.m. June 13, 7:00 a.m. – 7:30 p.m. June 14, 7:00 a.m. – 7:00 p.m. June 15, 7:00 a.m. – 1:00 p.m. MiCo – Milano Congressi South Wing, Level 0	"Spontaneous Pigmentation Spots are Signs of Successful RECELL® Therapy in Patients with Stable Large Area Vitiligo"	**Zhi-Fei Liu, et al** Peking Union Medical College Hospital Beijing, China

ABOUT AVITA MEDICAL LIMITED

AVITA Medical is a regenerative medicine company with a technology platform positioned to address unmet medical needs in burns, chronic wounds, and aesthetics indications. AVITA Medical's 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 REGENERATIVE EPIDERMAL SUSPENSION™ (RES™), 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 Medical's 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 7,000 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, acute wounds, scars and vitiligo. The RECELL System is TGA-registered in Australia and CFDA-cleared in China.

To learn more, visit 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.

FOR FURTHER INFORMATION:

U.S. Media **Sam Brown, Inc.** Christy Curran Phone +1-615.414.8668 christycurran@sambrown.com **O.U.S Media** **Monsoon Communications** Sarah Kemter Phone +61 (0)3 9620 3333 Mobile +61 (0)407 162 530 sarahk@monsoon.com.au	**Investors:** **Westwicke Partners** Caroline Corner Phone +1-415-202-5678 caroline.corner@westwicke.com

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