

**AVITA MEDICAL SUCCESSFULLY COMPLETES ISO QUALITY RECERTIFICATION AUDIT**

- **Receives ISO 9000, ISO 13485 and Med Device Directive Re-Certification**
- **Free to CE Mark all regenerative and respiratory products for sale in the European Union**

**Australia, 9 February 2012**—**Avita Medical Ltd.** (ASX: **AVH**), the regenerative medicine company, announced that, following the Quality Audit in November of 2011 by an independent, external auditor of the European Notified Body, the Company has received notification of successful audit results covering all applicable elements of ISO 9001:2008 Quality Management System Certification, ISO 13485:2003 Medical Devices Certification, Medical Devices Directive 93/42/EEC and Canadian Medical Devices Regulation Part for Avita's Class III Medical Devices. A rigorous and comprehensive recertification audit to all of the required elements is conducted only once every three years.

The audit addressed all phases of Avita Medical's Quality, Manufacturing and Management Systems and verified the controls on its portfolio of medical products including the Regenerative product, ReCell® Spray-On Skin®, as well as the company's respiratory products (Funhaler® and Breath-A-Tech®).

Successful ISO recertification is a requirement in order for Avita Medical to maintain its CE mark on its regenerative and respiratory products for sale within the European Union and Canada as well as a reference for registration in other countries, for instance, within the Middle East.

Mr William Marshall, Avita Medical's VP Operations and Regulatory said: "We are very pleased with this result. Over the last three years Avita has fully revamped and streamlined the company's Quality System and this successful audit demonstrates our continued commitment to the quality of our products and to our customers.

"Tight control of Quality and Manufacturing Systems are a prerequisite for efficient operations and are at the core of our Continuous Improvement model. The initiatives implemented have yielded significantly improved margins and reductions in operating costs."

**ABOUT AVITA MEDICAL LTD.**

Avita Medical ([www.avitamedical.com](http://www.avitamedical.com)) develops and distributes regenerative and tissue-engineered products for the treatment of a broad range of wounds, scars and skin defects. Using patented and proprietary tissue-culture, collection and application technology, the company is able to provide innovative treatment solutions derived from a patient's own skin. The company's lead product, ReCell Spray-On Skin, has been designed for use in a wide variety of burns, plastic, reconstructive and cosmetic procedures. ReCell is patented, CE-marked for Europe, TGA-registered in Australia, and SFDA-cleared in China. ReCell is not available for sale in the United States; in the U.S. ReCell is an investigational device limited by federal law to investigational use.

This news release may include forward-looking statements that involve risks and uncertainties. You can identify these statements by the use of words such as "anticipate", "estimate", "expect", "project", "potential", "intend", "plan", "believe", "target", "may", "assume" or similar expressions. These forward-looking statements speak only as at the date of this release and are based on management's expectations and beliefs concerning future events. Forward-looking statements are necessarily subject to risks, uncertainties and other factors, many of which are outside the control of Avita Medical that could cause actual results to differ materially from such statements. Avita Medical makes no undertaking to subsequently update or revise the forward-looking statements made in this release to reflect events or circumstances after the date of this release. This document is intended to provide background information only and does not purport to make any recommendation upon which you may reasonably rely without taking further and more specific advice.

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