



UNIVERSITY OF VIRGINIA HEALTH TO BECOME SECOND U.S. SITE FOR VISABL-AFL

15 December 2025 – Melbourne, Australia (**14 December 2025** – Minneapolis, MN United States) – **Imricor Medical Systems, Inc. (Company or Imricor) (ASX: IMR)** is excited to announce that the Institutional Review Board (**IRB**) of the University of Virginia (**UVA**) Health Medical Center has approved UVA Health's enrolment in Imricor's VISABL-AFL clinical trial.

Imricor personnel performed a Site Initiation Visit with five UVA Health physicians and UVA Health's Clinical Research Coordinator team on 11 December. Next steps include final approval of the agreed-upon site research contract, expected in the coming days, followed by site installation and clinical training, both planned for January after the holidays. Procedures are expected to commence in February.

The Principal Investigator (**PI**) for VISABL-AFL at UVA Health is Dr. Kenneth Bilchick, Electrophysiologist, Professor of Cardiology, and Director of Electrophysiology Research at UVA Health.

Dr. Bilchick commented: "The University of Virginia is pleased to be the 2nd site in the United States for the Imricor VISABL-AFL study. Interventional CMR, particularly for electrophysiology applications, promises to advance our therapeutic strategies for patients with atrial and ventricular arrhythmias by facilitating visualization of the catheters used for ablation simultaneously with real-time CMR imaging. We look forward to working with the other US and European VISABL-AFL sites to expand the role of MRI-guided electrophysiology ablations globally."

UVA Health is an academic health system associated with the University of Virginia in Charlottesville, Virginia, USA. The UVA Health University Medical Center includes a 600-plus-bed hospital and nationally ranked specialty programs. Newsweek and U.S. News & World Report consistently recognize it as the **top hospital in Virginia** and one of the **leading hospitals nationwide**.

Imricor's Chair and CEO, Steve Wedan, added: "The speed at which UVA Health has progressed from initial introduction to clearing IRB has been outstanding, and we are very excited to start procedures at a second US site. In addition, UVA Health is just the first of several US sites which are expected to join VISABL-AFL, helping to ensure that our enrolment and FDA processes remain on schedule."

ENDS

Authorised for release by Steve Wedan, Executive Chair, President, and CEO

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About Imricor

Imricor Medical Systems, Inc. (ASX:IMR) is striving to make interventional medical procedures better, safer, and more cost effective by making it possible for these procedures to be performed under real-time magnetic resonance imaging (MRI) guidance, rather than under x-ray fluoroscopy guidance, thus taking advantage of MRI's superior imaging capabilities.

Imricor's Products

Imricor is a pioneer and world leader in developing MRI-compatible products for cardiac catheter ablation procedures. The Company's products include capital equipment, such as the NorthStar® Mapping System and the Advantage-MR® EP Recorder/Stimulator. Single-use devices include a variety of ablation catheters, diagnostic catheters, steerable sheaths, and other tools used for cardiac ablations.

Imricor's products are approved in the European Union, the Kingdom of Saudi Arabia, and New Zealand. US FDA approval is in process, and further approvals in other geographies such as Australia are being planned.

Foreign Ownership Restrictions

Imricor's CHESS Depositary Interests (**CDIs**) are issued in reliance on the exemption from registration contained in Regulation S of the US Securities Act of 1933 (**Securities Act**) for offers which are made outside the US. Accordingly, the CDIs have not been, and will not be, registered under the Securities Act or the laws of any state or other jurisdiction in the US. As a result of relying on the Regulation S exemption, the CDIs are 'restricted securities' under Rule 144 of the Securities Act. This means that you are unable to sell the CDIs into the US or to a US person for the foreseeable future except in very limited circumstances after the expiration of a restricted period, unless the re-sale of the CDIs is registered under the Securities Act or an exemption is available. To enforce the above transfer restrictions, all CDIs issued bear a 'FOR US' designation on the Australian Securities Exchange (**ASX**). This designation restricts any CDIs from being sold on ASX to US persons. However, you are still able to freely transfer your CDIs on ASX to any person other than a US person. In addition, hedging transactions with regard to the CDIs may only be conducted in accordance with the Securities Act.

Forward-Looking Statements

This announcement contains or may contain forward-looking statements that are based on the Company's management's beliefs, assumptions and expectations and on information currently available to management. All statements that address operating performance, events or developments that we expect or anticipate will occur in the future are forward-looking statements. These include, without limitation, EU commercial market acceptance and EU sales of our product as well as our expectations with respect to our ability to develop and commercialise new products. Management believes that these forward-looking statements are reasonable when made. You should not place undue reliance on forward-looking statements because they speak only as of the date when made. Imricor does not assume any obligation to publicly update or revise any forward-looking statements, whether as a result of new information, future events or otherwise. Imricor may not actually achieve the plans, projections or expectations disclosed in forward-looking statements. Actual results, developments or events could differ materially from those disclosed in the forward-looking statements.

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