

Results of Special Meeting and PARADIGM Trial Update

MINNEAPOLIS, United States and BRISBANE, Australia 30 September 2025: Anteris Technologies Global Corp. (Anteris or the Company) (NASDAQ: AVR, ASX: AVR) a global structural heart company committed to designing, developing, and commercializing cutting-edge medical devices to restore healthy heart function, announces that on September 29, 2025, the Company held a Special Meeting of Stockholders (the **Special Meeting**) at which a quorum was present. The matters listed below were submitted to a vote of the Company’s stockholders at the Special Meeting through the solicitation of proxies. Detailed descriptions of the proposals are included in the Company’s definitive proxy statement on Schedule 14A filed with the Securities and Exchange Commission on August 18, 2025 (the “Proxy Statement”).

A total of 18,687,740 shares of the Company’s common stock were present at the Special Meeting in person, by virtual attendance, or by proxy, which represents approximately 51.8% of the shares of common stock outstanding as of August 11, 2025, which was the record date for the Special Meeting.

Proposal 1. The ASX Waiver Proposal

The Company’s stockholders approved ASX Limited's grant to the Company of a waiver from Australian Securities Exchange Listing Rule 7.1 to permit the Company to issue new securities without obtaining the approval of the Company's stockholders under ASX Listing Rule 7.1, and otherwise on the terms and conditions summarized in the Proxy Statement (the **ASX Waiver Proposal**). The final voting results are as follows:

Votes For	Votes Against	Abstentions
13,655,466	2,601,191	2,431,083

Proposal 2. The Adjournment Proposal

In connection with the Special Meeting, the Company’s stockholders approved the proposal to approve the adjournment of the Special Meeting, if necessary, to solicit additional proxies if there are not sufficient votes in favor of the ASX Waiver Proposal. No adjournment was necessary as Proposal 1 was approved. The final voting results are as follows:

Votes For	Votes Against	Abstentions
13,762,610	2,461,912	2,463,218

PARADIGM Trial Update

Patient enrolment in the Company’s global pivotal registration trial (the **PARADIGM Trial**), initially expected in the third quarter 2025, is now anticipated to commence in the fourth quarter 2025.



For personal use only

Following the Investigational Device Exemption (IDE) review, the Federal Drug Administration (FDA) requested additional information, primarily related to bench testing, which the Company has now addressed in a formal response. The IDE submission incorporates both the clinical study protocol and the underlying technology package, which demonstrates the DurAVR® THV System has been appropriately developed and tested for use in the global pivotal trial.

In parallel, regulatory submissions to competent authorities in Europe are underway, highlighting the Company's commitment to advancing the PARADIGM Trial across multiple regions.

About Anteris

Anteris Technologies Global Corp. (NASDAQ: AVR, ASX: AVR) is a global structural heart company committed to designing, developing, and commercializing cutting-edge medical devices to restore healthy heart function. Founded in Australia, with a significant presence in Minneapolis, USA, Anteris is a science-driven company with an experienced team of multidisciplinary professionals delivering restorative solutions to structural heart disease patients.

Anteris' lead product, the DurAVR® Transcatheter Heart Valve (THV), was designed in partnership with the world's leading interventional cardiologists and cardiac surgeons to treat aortic stenosis – a potentially life-threatening condition resulting from the narrowing of the aortic valve. The balloon-expandable DurAVR® THV is the first biomimetic valve, which is shaped to mimic the performance of a healthy human aortic valve and aims to replicate normal aortic blood flow. DurAVR® THV is made using a single piece of molded ADAPT® tissue, Anteris' patented anti-calcification tissue technology. ADAPT® tissue, which is FDA-cleared, has been used clinically for over 10 years and distributed for use in over 55,000 patients worldwide. The DurAVR® THV System is comprised of the DurAVR® valve, the ADAPT® tissue, and the balloon-expandable ComASUR® Delivery System.

Authorisation and Additional information

This announcement was authorised for release on the ASX by the Chief Executive Officer, Wayne Paterson.

For more information:

Investor Relations

investors@anteristech.com
Debbie Ormsby
Anteris Technologies Global Corp.
+61 1300 550 310 | +61 7 3152 3200

Investor Relations (US)

mchatterjee@bplifescience.com
Malini Chatterjee, Ph.D.
Blueprint Life Science Group
+1 917 330 4269

Website	www.anteristech.com
X	@AnterisTech
LinkedIn	https://www.linkedin.com/company/anteristech

