Anteris Advances Global PARADIGM Expansion with French Regulatory Clearance

MINNEAPOLIS, United States and **BRISBANE, Australia 9 June 2026: Anteris Technologies Global Corp.** ("Anteris" or the "Company") (NASDAQ: AVR, ASX: AVR) announced today that it has received full regulatory clearance from the French National Agency for Medicines and Health Products Safety (ANSM) for the DurAVR® Transcatheter Heart Valve (THV) global pivotal trial in patients with severe calcific aortic stenosis (the "PARADIGM Trial"). This authorisation marks a significant milestone in the Company's clinical and regulatory strategy and enables patient recruitment to commence at leading centres in France.

"Securing French regulatory clearance is an important step in the execution of the PARADIGM Trial," said Wayne Paterson, Vice Chairman and Chief Executive Officer of Anteris. "France represents a strategically important market with highly experienced centres and investigators, reinforcing the quality and conduct of the study".

The Company continues to advance its global clinical strategy, with ongoing site activations and patient enrolment across the United States and Europe.

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About the PARADIGM Trial

The PARADIGM Trial is a prospective randomized controlled trial which will evaluate the safety and effectiveness of the DurAVR® Transcatheter Heart Valve compared to commercially available transcatheter aortic valve replacements (TAVRs).

This head-to-head study will enrol approximately 1,000 patients in the 'All Comers Randomized Cohort' with 1:1 randomization of patients who will receive either the DurAVR® THV or TAVR using commercially available and approved THVs. The PARADIGM Trial will assess non-inferiority on a primary composite endpoint of all-cause mortality, all stroke and cardiovascular hospitalization at one year post procedure.

For further information, please refer to ClinicalTrials.gov NCT07194265.

About Anteris

Anteris Technologies Global Corp. (NASDAQ: AVR, ASX: AVR) is a global healthcare 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® THV, was designed in collaboration 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.

Forward-Looking Statements

This announcement contains forward-looking statements, including statements regarding the expectation that patient recruitment will commence in France. Forward-looking statements include all statements that are not historical facts. Forward-looking statements generally are identified by the words "believe," "project," "expect," "anticipate," "estimate," "intend," "budget," "target," "aim," "strategy," "plan," "guidance," "outlook," "may," "should," "could," "will," "would," "will be," "will continue," "will likely result" and similar expressions, although not all forward-looking statements contain these identifying words. These forward-looking statements are subject to a number of risks, uncertainties, and assumptions, including those described under "Risk Factors" in Anteris' Annual Report on Form 10-K for the fiscal period ended December 31, 2025 that was filed with the Securities and Exchange Commission and ASX. Actual future events may vary from these forward-looking statements and readers are cautioned not to put undue reliance on forward-looking statements. Other than as required by law, Anteris gives no representation or guarantee that the occurrence of any of the events or circumstances expressed or implied in these statements will occur. In addition, except as required by law, Anteris does not assume any obligation to update any of these forward-looking statements to conform these statements to actual results or revised expectations.

Authorisation and Additional information

This announcement was authorised for release on the ASX by the Board of Directors.

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