UNITED STATES
SECURITIES AND EXCHANGE COMMISSION
Washington, D.C. 20549

FORM 8-K

CURRENT REPORT
Pursuant to Section 13 OR 15(d)
of The Securities Exchange Act of 1934

Date of Report (Date of earliest event reported): November 10, 2026

Anteris Technologies Global Corp.

(Exact name of registrant as specified in its charter)

Delaware	**001-42437**	**99-1407174**
(State or Other Jurisdiction of Incorporation)	**(Commission File Number)**	**(I.R.S. Employer Identification No.)**

Toowong Tower, Level 3, Suite 302 **9 Sherwood Road** **Toowong, QLD** **Australia**	**4066**
(Address of Principal Executive Offices)	**(Zip Code)**

Registrant's telephone number, including area code: +61 7 3152 3200

Not Applicable
(Former name or former address, if changed since last report)

Check the appropriate box below if the Form 8-K filing is intended to simultaneously satisfy the filing obligation of the registrant under any of the following provisions:

☐ Written communications pursuant to Rule 425 under the Securities Act (17 CFR 230.425)

☐ Soliciting material pursuant to Rule 14a-12 under the Exchange Act (17 CFR 240.14a-12)

☐ Pre-commencement communications pursuant to Rule 14d-2(b) under the Exchange Act (17 CFR 240.14d-2(b))

☐ Pre-commencement communications pursuant to Rule 13e-4(c) under the Exchange Act (17 CFR 240.13e-4(c))

Securities registered pursuant to Section 12(b) of the Act:

Title of each class	**Trading Symbol(s)**	**Name of each exchange on which registered**
Common Stock, par value $0.0001 per share	AVR	The Nasdaq Global Market

Indicate by check mark whether the registrant is an emerging growth company as defined in Rule 405 of the Securities Act of 1933 (§230.405 of this chapter) or Rule 12b-2 of the Securities Exchange Act of 1934 (§240.12b-2 of this chapter).

Emerging growth company ☒

If an emerging growth company, indicate by check mark if the registrant has elected not to use the extended transition period for complying with any new or revised financial accounting standards provided pursuant to Section 13(a) of the Exchange Act. ☐

Item 2.02. Results of Operations and Financial Condition.

On November 10, 2026 USA ET (November 11, 2026 AEST), Anteris Technologies Global Corp. (the "Company") (i) filed the Form 10-Q for the quarter ended September 30, 2026 with a cover page (the "Results Announcement") with the Australian Securities Exchange ("ASX"); and (ii) issued an ASX Announcement regarding the Company's financial results for the quarter ended September 30, 2026, both of which include unaudited and other historical financial information for the period ended September 30, 2026. The Results Announcement has been prepared for the purpose of complying with the reporting requirements of the ASX. Copies of the Results Announcement and the ASX Announcement are attached as Exhibits 99.1 and 99.2, respectively, to this Current Report on Form 8-K.

The information in this Item 2.02, including Exhibits 99.1 and 99.2 attached hereto, is being furnished and shall not be deemed to be filed for the purposes of Section 18 of the Securities Exchange Act of 1934 (the "Exchange Act"), or incorporated by reference into any filing under the Securities Act of 1933 or the Exchange Act, unless such subsequent filing specifically references this Current Report on Form 8-K.

Item 9.01. Financial Statements and Exhibits.

(d) Exhibits.

The following exhibits are filed with this Current Report on Form 8-K:

Exhibit No.	Description
99.1	Results Announcement, as filed with the ASX, for the quarter ended September 30, 2026
99.2	ASX Announcement regarding financial results for the period ended September 30, 2026
104	Cover Page Interactive Data File (embedded within the Inline XBRL document)

SIGNATURES

Pursuant to the requirements of the Securities Exchange Act of 1934, the registrant has duly caused this report to be signed on its behalf by the undersigned hereunto duly authorized.

Anteris Technologies Global Corp.

Date: November 10, 2026

By: /s/ Wayne Paterson
Name: Wayne Paterson
Title: Vice Chairman and Chief Executive Officer

Results for announcement to the market

Name of entity: Anteris Technologies Global Corp. ("ATGC")
ARBN: 677 960 235
Reporting period: For the quarter ended September 30, 2026

The attached Form 10-Q *Quarterly Report* for the quarter ended September 30, 2026 (the "Form 10-Q") has been filed with the U.S. Securities and Exchange Commission ("SEC"). It includes the condensed consolidated financial statements which have been prepared in accordance with generally accepted accounting principles in the United States of America ("U.S. GAAP") and are denominated in U.S. dollars.

The following supplementary information is provided in connection with the Form 10-Q to satisfy the Company's ongoing disclosure obligations under the ASX Listing Rules. The information has been prepared in accordance with the waiver conditions granted by the ASX in relation to 4.7C for quarterly activity reports, and should be read in conjunction with the Form 10-Q.

Details of business activities during the quarter

Refer to the Form 10-Q and the "Anteris Announces Results for the Third Quarter of 2026" announcement lodged with the ASX on November 11, 2026.

Aggregate amount of payments to related parties and their associates

During the third quarter of 2026, the aggregate amount of payments to related parties and their associates was US$576 thousand. The quarterly amount includes fees paid to non-executive directors, as well as salary payments to the CEO, President and CFO. These payments were included in cash flows from operating activities.

There were no payments to related parties or their associates included in cash flows from investing activities.

ENDS

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.

860 Blue Gentian Road,
Suite 340
Eagan, MN, 55121
United States
T: +1 651 493 0606
info.us@anteristech.com

Anteris Technologies Global Corp.
BRISBANE | MINNEAPOLIS | GENEVA | MALAGA
anteristech.com

Toowong Tower, Level 3, Suite 302
9 Sherwood Road, Toowong
QLD 4066, Australia
T: +61 1300 550 310
info.au@anteristech.com
ARBN: 677 960 235

Forward-Looking Statements

This announcement contains forward-looking statements. 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 and the Form 10-Q, each filed with the SEC 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.

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

Anteris Technologies Announces Results for the Third Quarter of 2026

MINNEAPOLIS, United States and **BRISBANE, Australia November 11, 2026 (AEDT): Anteris Technologies Global Corp.** ("Anteris" or the "Company") (NASDAQ: AVR, ASX: AVR), a global healthcare company committed to designing, developing, and commercializing cutting-edge medical devices to restore healthy heart function, today reported financial results for the quarter ended September 30, 2026, and provided a corporate update.

Q3 2026 Highlights

- Continued execution of the PARADIGM Trial, with enrollment progressing across active sites in the U.S., Canada, Denmark, the Netherlands and France. Subsequent to quarter-end, the trial received regulatory clearance from Germany's Federal Institute for Drugs and Medical Devices (BfArM) and advanced in Australia following HREC approval and TGA authorization under the Clinical Trial Notification pathway.

- Assumed occupancy of the Northcross facility in Minnesota, providing additional capacity to scale production of the DurAVR® THVs and ComASUR® Delivery System, in preparation for commercialization.

- Appointed Brent Moen as Chief Financial Officer, effective September 11, to support the Company's financial and governance priorities as it advances the DurAVR THV toward commercialization.

"Q3 saw continued progress on multiple fronts. The PARADIGM Trial recruitment is advancing along with capacity expansion in line with our clinical and commercial plan. The Company also brought in some key hires to strengthen our operating model and future capabilities," said Wayne Paterson, Vice Chairman and Chief Executive Officer of Anteris.

Business & Operations

During the quarter, the Company continued to execute the global pivotal PARADIGM Trial across active sites in the United States, Canada and Europe.

The Company increased site activation in the United States and expanded the PARADIGM Trial into Canada and France, where sites are active and enrolling, following completion of key start-up activities, including site qualification, investigator training and operational readiness. Subsequent to quarter end, the trial received regulatory clearance from the German Federal Institute for Drugs and Medical Devices (BfArM), supporting its expansion into Germany. In Australia, the trial received Human Research Ethics Committee (HREC) approval and authorization from the Therapeutic Goods Administration (TGA) under the Clinical Trial Notification (CTN) pathway, further extending the trial's geographic reach.

On September 1, the Company assumed occupancy as the primary tenant of the Northcross facility in Brooklyn Park, Minnesota. The facility provides additional capacity to scale production of the DurAVR THVs, the ComASUR Delivery System and related components in preparation for commercialization. The ComASUR Delivery System is designed to support controlled deployment and accurate placement of the balloon-expandable DurAVR THV, a new class of biomimetic transcatheter aortic valve.

Financial Results

The financial results for Anteris for the quarter ended September 30, 2026, are presented below. All amounts in $ refer to U.S. dollars.

The Company's net operating cash outflows for the three months ended September 30, 2026, were $22.9 million, primarily attributable to clinical, regulatory and manufacturing requirements to support the ongoing PARADIGM Trial. Operating expenditures reflected the phased execution of the clinical program during the quarter, including the timing of U.S. site activation activities following receipt of the CMS coverage determination in April 2026. R&D expenses of $21.3 million were driven by the scaling of manufacturing and quality capabilities, including process development and validation activities and expanded headcount, together with PARADIGM trial related activities, including clinical costs associated with patient enrollment and the scaling of our field-based clinical team. These costs were partly offset by reduced DurAVR THV product research costs.

Please see the detailed financial information contained in Anteris' Quarterly Report on Form 10-Q for the quarter ended September 30, 2026.

ENDS

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 ("THV") compared to commercially available transcatheter aortic valve replacements (TAVRs).

This head-to-head study will enroll 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.

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DurAVR THV is intended for INVESTIGATIONAL USE ONLY. Not available for commercial use or sale worldwide. CAUTION—Investigational device. Limited by Federal (or United States) law to investigational use.

Forward-Looking Statements

This announcement contains forward-looking statements, including, but not limited to, statements regarding the PARADIGM Trial, clinical development timelines and potential commercialization. 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.

Authorization and Additional Information

This announcement was authorized for release on the ASX by Wayne Paterson, Vice Chairman and Chief Executive Officer.

For more information:

Global 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