

## **ACURATE neo2™ Aortic Valve System data presented at late-breaking clinical trial session at TCT 2024**

MARLBOROUGH, Mass. and WASHINGTON, Oct. 30, 2024 [/PRNewswire/](#) -- Boston Scientific Corporation (NYSE: BSX) today announced the results of the primary endpoint of the ACURATE IDE clinical trial, which evaluated the ACURATE neo2™ Aortic Valve System in the treatment of patients with severe, symptomatic aortic stenosis at low, intermediate, high and extreme risk of open-heart surgery. The data were presented as a late-breaking clinical trial at Transcatheter Cardiovascular Therapeutics® (TCT®) 2024, the annual scientific symposium of the Cardiovascular Research Foundation® (CRF®).

This randomized trial evaluated the ACURATE neo2 valve, the company's second-generation transcatheter aortic valve replacement (TAVR) technology, versus a pre-defined control valve, either the commercially available SAPIEN™ valve or the Evolut™ transcatheter aortic valve system, selected at the discretion of the implanting physician. The composite rate of all-cause mortality, stroke or rehospitalization at one year was 16.16% in the ACURATE neo2 arm and 9.53% in the control arm, not meeting the prespecified criterion for non-inferiority (posterior probability for non-inferiority was 77.9%, which is lower than the non-inferiority test threshold of 97.5%).

"This trial is the largest randomized comparison of TAVR platforms and was conducted over four years, including during a global pandemic that introduced complexities for enrollment," said Michael Reardon, M.D., professor of cardiothoracic surgery at Houston Methodist DeBakey Heart & Vascular Center and co-principal investigator of the ACURATE IDE trial. "These data add to the breadth of clinical knowledge of the ACURATE valve platform and provide compelling insights on the importance of procedural optimization that will be beneficial for TAVR moving forward."

Also presented in the late-breaking session were data from a post-hoc analysis to identify and evaluate expanded and under-expanded ACURATE neo2 valve frames within the ACURATE IDE trial. The company performed a review of key procedural factors, including pre- and post-dilation, which helps to prepare for, enable and confirm proper valve expansion during the procedure. Following this review, an assessment of implant quality for the ACURATE neo2 valve was initiated and highlighted that approximately 20% of the valves were under-expanded. Data from the assessment also demonstrated that the rate for death, stroke or rehospitalization at one year was similar between the ACURATE neo2 expanded group and the control group.

"The data presented today give clinicians a greater understanding of the impact of procedural optimization as the TAVR space continues to rapidly evolve," said Janar Sathanathan, M.D., chief medical officer, Interventional Cardiology Therapies, Boston Scientific. "We believe the findings from the ACURATE IDE post-hoc analysis and implementation of steps to mitigate valve under-expansion may improve outcomes for the ACURATE valve and have important implications on other commercially available TAVR valves, and we look forward to studying these improved techniques in future trials of the device."

Boston Scientific continues to work closely with the U.S. Food and Drug Administration on the regulatory strategy for approval of the ACURATE valve platform in the U.S.

More information on the ACURATE IDE trial is available [here](#).

\*In Europe, the ACURATE neo2™ Aortic Valve System and the ACURATE Prime™ Aortic Valve System are CE-marked. In the USA, the ACURATE neo2 Aortic Valve System and the ACURATE Prime Aortic Valve System are investigational devices and are restricted under federal law to investigational use only. Not available for sale.

\*\*Dr. Michael Reardon is a paid consultant of Boston Scientific Corporation. He has not been compensated in connection with this press release.

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This press release contains forward-looking statements within the meaning of Section 27A of the Securities Act of 1933 and Section 21E of the Securities Exchange Act of 1934. Forward-looking statements may be identified by words like "anticipate," "expect," "project," "believe," "plan," "estimate," "intend" and similar words. These forward-looking statements are based on our beliefs, assumptions and estimates using information available to us at the time and are not intended to be guarantees of future events or performance. These forward-looking statements include, among other things, statements regarding our business plans and product performance and impact, clinical trials, and new and anticipated product approvals and launches. If our

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Factors that may cause such differences include, among other things: future economic, competitive, reimbursement and regulatory conditions; geopolitical events; manufacturing, distribution and supply chain disruptions and cost increases; disruptions caused by cybersecurity events; disruptions caused by public health emergencies or extreme weather or other climate change-related events; labor shortages and increases in labor costs; variations in outcomes of ongoing and future clinical trials and market studies; new product introductions; expected procedural volumes; the closing and integration of acquisitions; demographic trends; intellectual property; litigation; financial market conditions; and future business decisions made by us and our competitors. All of these factors are difficult or impossible to predict accurately and many of them are beyond our control. For a further list and description of these and other important risks and uncertainties that may affect our future operations, see Part I, Item 1A – Risk Factors in our most recent Annual Report on Form 10-K filed with the Securities and Exchange Commission, which we may update in Part II, Item 1A – Risk Factors in Quarterly Reports on Form 10-Q we have filed or will file hereafter. We disclaim any intention or obligation to publicly update or revise any forward-looking statements to reflect any change in our expectations or in events, conditions or circumstances on which those expectations may be based, or that may affect the likelihood that actual results will differ from those contained in the forward-looking statements, except as required by law. This cautionary statement is applicable to all forward-looking statements contained in this document.

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Additional assets available online: [Photos \(1\)](#)

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