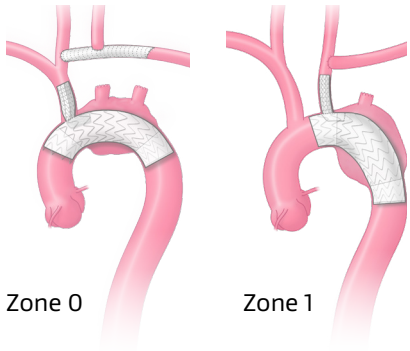
GORE® TAG®
Thoracic Branch Endoprosthesis

Evaluation of the GORE® TAG® Thoracic Branch Endoprosthesis (TBE) in the Treatment of Lesions of the Aortic Arch and Descending Thoracic Aorta (Zone 0/1)

Non–randomized, multicenter study

- Subjects with Zone 0/1 Aortic aneurysm or Zone 0/1 non-aneurysm aortic lesions, including dissection and other isolated lesion types, treated with the GORE® TAG® Thoracic Branch Endoprosthesis.
- Pivotal study SSB 11-02.
- ClinicalTrials.gov Identifier: NCT02777528.
- Clinical protocol is approved for 100 subjects across three cohorts (aneurysm, dissection and isolated lesions) at up to 40 sites.



CAUTION: Investigational device. Limited by United States law to investigational use in Zone 0/1. Approved for use in Zone 2 by the U.S. Food and Drug Administration (FDA).

U.S. clinical sites

Baylor College of Medicine – Houston	Cardiothoracic and Vascular Surgeons – Austin	Cardiovascular Surgery Clinic
Carolinas HealthCare Systems	Cedar–Sinai Medical Center	Cleveland Clinic
Cooper University Hospital	Dartmouth – Hitchcock Medical Center	Duke University Medical Center
Emory University	Hartford Hospital	Heart Hospital at Baylor Plano
Hospital of the University of Pennsylvania	Houston Methodist Hospital	Keck Medical Center of University of Southern California (USC)
Leland Stanford Junior University	Massachusetts General Hospital	Mayo Clinic – Rochester
MedStar Washington Hospital Center	Memorial Hermann Health System	Northwestern University
Oregon Health & Science University	Research Foundation SUNY Buffalo	Sentara Medical Group
St. Vincent Medical Group, Inc.	University of Baltimore Maryland	University of Florida – Gainesville
University of Michigan	University of Pittsburgh Medical Center	University of South Florida
University of Washington	University of Wisconsin System	Vanderbilt University Medical Center
Washington University School of Medicine – St. Louis	William Beaumont Hospital	

Products listed may not be available in all markets.

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