

## Real-world data

### GORE® TAG® Conformable Thoracic Stent Graft with ACTIVE CONTROL System



**SURPASS\* is an observational, prospective, single-arm post-market registry. 20 European sites in seven countries are included.**

# ZERO

- Type Ia and Type III endoleaks
- Fractures
- Device compressions
- Ruptures

## 100%

Successful deployment

## 98.4%

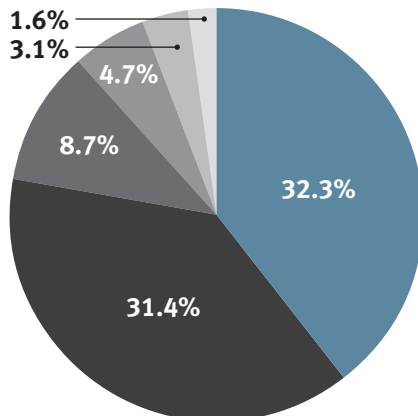
With no device-related issues

## 97.2%

Freedom from serious access complications

#### Aortic pathology treated:

- Treatment of Type B dissection (complicated / uncomplicated)
- Descending thoracic aortic aneurysms (Including ruptures)
- Penetrating aortic ulcers
- Traumatic aortic transection
- Intramural hematoma
- Pseudoaneurysm



## 98.4%

Reported that proximal wall apposition was acceptable at procedural completion

## 1.38

Devices per procedure

## 92.9%

No rapid pacing used

\* Rates are based on physician experience as reported for 127 subjects in Europe within a 30 day follow-up period.  
European-Post Approval Registry: Observational Registry Characterizing the Performance and Feature Use of the GORE® TAG® Conformable Thoracic Stent Graft Featuring ACTIVE CONTROL System. (data on file 2017; W.L. Gore & Associates, Inc; Flagstaff, AZ.)

**INDICATIONS FOR USE IN THE U.S.:** The GORE® TAG® Thoracic Stent Graft is intended for endovascular repair of all lesions of the descending thoracic aorta, including: Isolated lesions in patients who have appropriate anatomy, including: adequate iliac / femoral access, aortic inner diameter in the range of 16–42 mm,  $\geq 20$  mm non-aneurysmal aorta proximal and distal to the lesion; Type B dissections in patients who have appropriate anatomy, including: adequate iliac / femoral access,  $\geq 20$  mm landing zone proximal to the primary entry tear; proximal extent of the landing zone must not be dissected, diameter at proximal extent of proximal landing zone in the range of 16–42 mm. **CONTRAINDICATIONS:** Patients with known sensitivities or allergies to the device materials; patients who have a condition that threatens to infect the graft. Refer to *Instructions for Use* at [goremedical.com](http://goremedical.com) for a complete description of all warnings, precautions, and adverse events. **INDICATIONS FOR USE UNDER CE MARK:** The GORE® TAG® Thoracic Stent Graft is indicated for endovascular repair of the descending thoracic aorta. **CONTRAINDICATIONS:** Patients with known sensitivities or allergies to the device materials; patients with a systemic infection who may be at increased risk of endovascular graft infection. Refer to *Instructions for Use* at [goremedical.com](http://goremedical.com) for a complete description of all warnings, precautions, and adverse events. Rx Only

Products listed may not be available in all markets.

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