Designed For Conformability
Building Confidence in the Treatment of Type B Dissection

Worldwide data supports performance in the treatment of all Type B dissections
Gore GREAT Registry provides worldwide experience and outcomes in real-world dissection use.

**GREAT Dissection Experience**

### Dissection Sub-Types Treated
- Acute Complicated: 39%
- Acute Uncomplicated: 23%
- Chronic: 38%

### Proximal Landing Zone
- Zone 0: 11%
- Zone 1: 4%
- Zone 2: 5%
- Zone 3: 11%
- Zone 4: 30%

### Adverse Event GREAT Data
- Dissection-Related Survival: 96.4%
- Procedural Survival: 100%
- RTAD: 0.6%
- Stroke: 1.2%
- Type IA Endoleak: 0%
- Device Compressions: 0%
- Paraparesis / Paralysis: 1.2%
- Conversion: 0.6%

1. The GREAT Registry is a prospective, observational, multicenter registry to actively track Gore commercial aortic endovascular device performance and associated patient outcomes in global markets with 10 years of follow-up.
2. The GORE® TAG® Thoracic Endoprosthesis is not indicated for the treatment of Zone 0 and Zone 1.
3. Includes all Type B dissections, events occurring within 12 months post-procedure.

INDICATIONS FOR USE IN THE US: The GORE® TAG® Thoracic Endoprosthesis 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, ≥ 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, ≥ 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 for a complete description of all warnings, precautions, and adverse events.

INDICATIONS FOR USE UNDER CE MARK: The GORE® TAG® Thoracic Endoprosthesis 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 for a complete description of all warnings, precautions, and adverse events.

products listed may not be available in all markets.

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To view further dissection information, visit goremedical.com/aortic/tevar