Trust is Earned

125,000
Devices Distributed

Proven long-term outcomes across all etiologies through five years

The Conformable GORE® TAG® Device has demonstrated long-term freedom from device-related reintervention (93.1%) and low complication rates (zero migrations, fractures, or compressions).

Aneurysm, Transection, All Type B Dissection

THROUGH FIVE-YEAR FOLLOW-UP*	(N = 217)				
Procedural survival	100%	Spinal cord ischemia	3.7%	Fracture	0%
30-day survival	95.4%	Paraplegia	0.5%	Compression	0%
Freedom from device-related reintervention	93.1%	Aortic rupture associated with treated area	1.8%	Migration	0%

* Consolidated outcomes following 5 years of follow-up in TAG 08-01, TAG 08-02, and TAG 08-03 clinical studies.

CONFORMABLE

GORE TAG THORACIC ENDOPROSTHESIS

Continuing with a legacy of firsts, the GORE® TAG® Device family ***was the first*** to reach 100,000 devices distributed worldwide. The Conformable GORE® TAG® Device has ***proven*** to be a safe, effective, and durable solution, earning the trust of physicians worldwide.

Gore TEVAR Leadership

1st approved device	**1st** approved device	**20** year
in the U.S., Europe, and Japan	for endovascular treatment of aneurysm, transection, and Type B dissection	clinical history

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. ℞ Only

Products listed may not be available in all markets.

GORE
Creative Technologies Worldwide

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