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

New levels of control delivering intraoperative TEVAR precision for hospital efficiency

Hospitals can count on efficient and reliable TEVAR outcomes even in challenging patient anatomies

Low rates of reintervention (3%)[1]

- Conformable stent graft adapts to the patient anatomy
- Built on the established success of the Conformable GORE® TAG® Device, one of the most trusted TEVAR devices on the market

Helps reduce the potential for complications

- New delivery system offers staged, controlled deployment

Value through efficiency

- Stock fewer devices while treating a broad range of patients
- Designed to reduce procedure time through intraoperative accuracy

For information on how this can impact your patient outcomes contact your local sales associate

goremedical.com/predictable

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1. W. L. Gore & Associates. 'GREAT' Global Registry for Endovascular Aortic Treatment - Outcomes Evaluation. Bethesda, MD: National Library of Medicine; 2012. Available from: https://clinicaltrials.gov/ct2/show/NCT01658787. NLM Identifier: NCT01658787. Published August 3, 2012. Updated: October 27, 2016. Accessed: Accessed June 22, 2017

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Catalogue number	Intended aortic diameter (mm)	Proximal diameter (mm)	Distal diameter (mm)	Endoprosthesis length (cm)	Recommended GORE® Dryseal Flex Introducer Sheath size (fr)	Oversizing range (%)	Partially uncovered stent length (mm)
TGM212110	16–19.5	21	21	10	18	8–31	3
TGM262110	19.5–24 / 16–19.5	26	21	10	20	8–33	4
TGM262610	19.5–24	26	26	10	20	8–33	4
TGM282810	22–26	28	28	10	20	8–27	4
TGM282815	22–26	28	28	15	20	8–27	4
TGMR312610	24–29 / 19.5–24	31	26	10	20	7–33	4
TGMR313110	24–29	31	31	10	20	7–29	4
TGMR313115	24–29	31	31	15	20	7–29	4
TGMR313120	24–29	31	31	20	20	7–29	4
TGM343410	27–32	34	34	10	22	6–26	5
TGM343415	27–32	34	34	15	22	6–26	5
TGM343420	27–32	34	34	20	22	6–26	5
TGMR373710	29–34	37	37	10	22	9–28	5
TGMR373715	29–34	37	37	15	22	9–28	5
TGMR373720	29–34	37	37	20	22	9–28	5
TGMR404010	31–37	40	40	10	22	8–29	6
TGMR404015	31–37	40	40	15	22	8–29	6
TGMR404020	31–37	40	40	20	22	8–29	6
TGM454510	34–42	45	45	10	24	7–32	6.5
TGM454515	34–42	45	45	15	24	7–32	6.5
TGM454520	34–42	45	45	20	24	7–32	6.5

For Europe, Australia, and New Zealand, add E at the end of the catalogue number.

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INDICATIONS FOR USE IN THE U.S.: The GORE® TAG® Conformable 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, ≥ 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® Conformable 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 for a complete description of all warnings, precautions, and adverse events. ℞ Only

Products listed may not be available in all markets.