

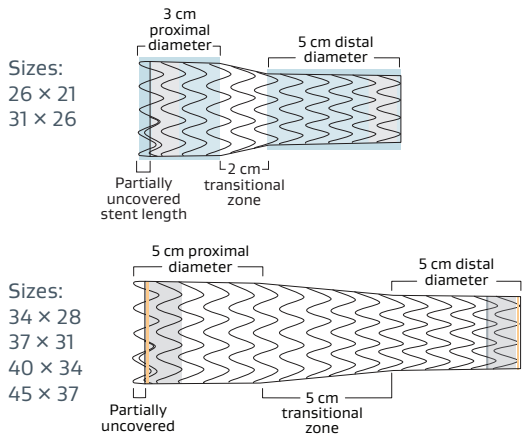
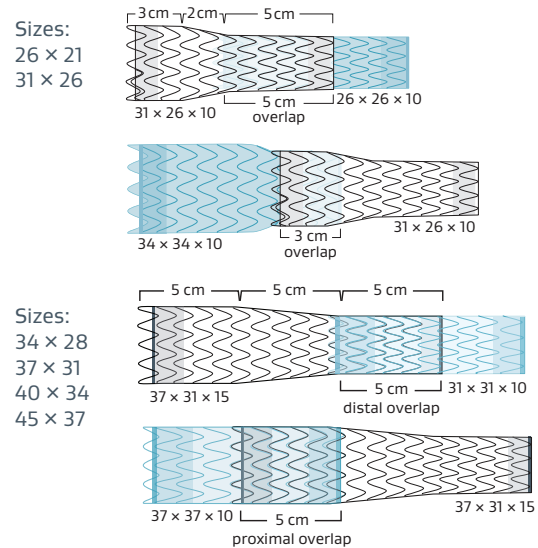
Using multiple devices

- Overlap ≥ 3 cm if second device is 1 or 2 sizes larger
- Overlap ≥ 5 cm if second device is the same size

Tapered devices

- The partially uncovered stents are part of the 2 cm proximal landing zone for straight and tapered devices

Overlapping a tapered device



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

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)
Straight Devices							
TGM212110	16-19.5	21	21	10	18	8-31	3
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
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
Tapered Devices							
TGM262110	19.5-24/16-19.5	26	21	10	20	8-33	4
TGMR312610	24-29/19.5-24	31	26	10	20	7-33	4
TGM342815	27-32/22-26	34	28	15	22	6-27	5
TGMR373115	29-34/24-29	37	31	15	22	7-29	5
TGMR403415	31-37/27-32	40	34	15	22	6-29	6
TGM453715	34-42/29-34	45	37	15	24	7-32	6.5

Required ancillary devices

GORE® DRYSEAL Flex Introducer Sheath

Catalogue number ^a	Sheath size (Fr)	Minimum sheath ID (mm)	Nominal sheath OD (mm)
GDSF1133	11	3.3	4.0
GDSF1833	18	6.0	6.7
GDSF1865	18	6.0	6.7
GDSF2033	20	6.7	7.5
GDSF2065	20	6.7	7.5
GDSF2233	22	7.3	8.2
GDSF2265	22	7.3	8.2
GDSF2433	24	8.0	8.8
GDSF2465	24	8.0	8.8
GDSF2633	26	8.7	9.5
GDSF2665	26	8.7	9.5

a. GDSF catalogue numbers are not available in all regions; use DSF catalogue numbers instead where appropriate.

 Consult Instructions for Use at eifu.goremedical.com

Refer to *Instructions for Use* at eifu.goremedical.com for a complete description of all applicable indications, warnings, precautions and contraindications for the markets where this product is available. Rx only

INDICATIONS FOR USE: 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.

Products listed may not be available in all markets.

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GORE® Tri-Lobe Balloon Catheter

	Catalogue number ^b	Inner vessel diameter (mm)
US:	TBCM1634	16-32
	TBCL2648	26-48
	Catalogue number ^b	Inner vessel diameter (mm)
Canada:	BCM1634	16-32
	BCL2648	26-48

b. Products listed may not be available in all markets. For accounts where TBCM1634 and TBCL2648 are not available, BCM1634 and BCL2645 may be substituted.

Other materials required for device placement

- 0.035" (0.89 mm) stiff guidewire, 260 cm or longer
- Heparin and heparinized saline solution
- Contrast medium
- Sterile syringes
- Three-way stopcock
- Appropriate diagnostic catheters and accessories

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