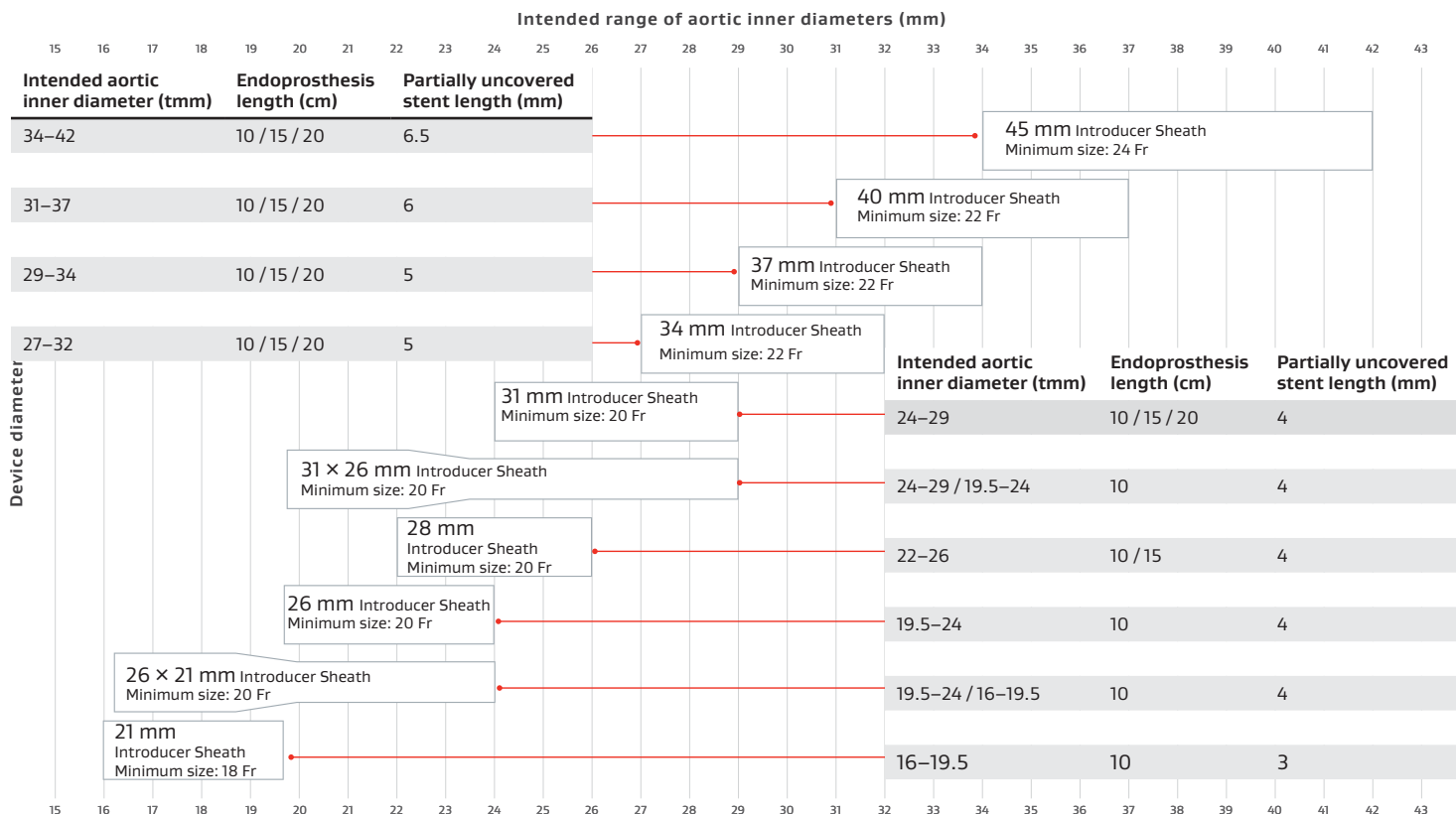


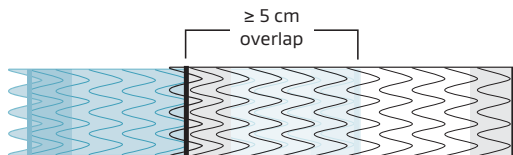


SIZING GUIDE

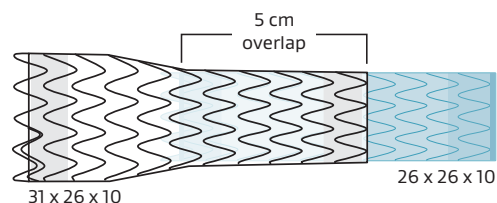


Using multiple devices

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

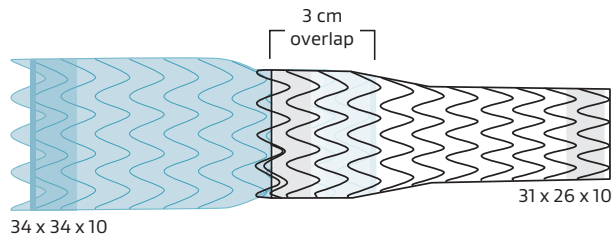
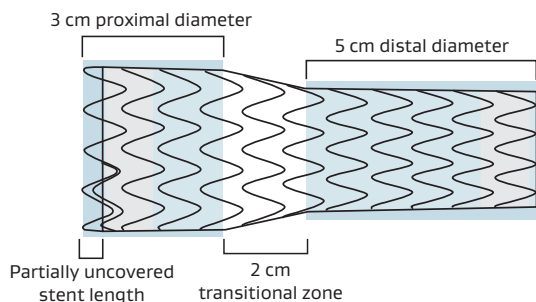


Overlapping a tapered device



Tapered devices

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



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)
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

Required ancillary devices

GORE® DRYSEAL Flex Introducer Sheath

Catalogue number	Sheath size (Fr)	Minimum sheath ID (mm)	Nominal sheath OD (mm)
DSFI833	18	6.0	6.7
DSF2033	20	6.7	7.5
DSF2065	20	6.7	7.5
DSF2233	22	7.3	8.2
DSF2265	22	7.3	8.2
DSF2433	24	8.0	8.8
DSF2465	24	8.0	8.8
DSF2633	26	8.7	9.5
DSF2665	26	8.7	9.5

GORE® Tri-Lobe Balloon Catheter

Catalogue number	Inner Vessel Diameter (mm)
BCM1634	16–32
BCL2645	26–42

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

 Consult Instructions for Use
eifu.goremedical.com

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 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 UNDER CE MARK: The GORE® TAG® Conformable Thoracic Stent Graft is indicated for endovascular repair of all lesions of the descending thoracic aorta, including isolated lesions, such as aneurysm and traumatic transection, and Type B dissections. **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 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

Products listed may not be available in all markets.

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