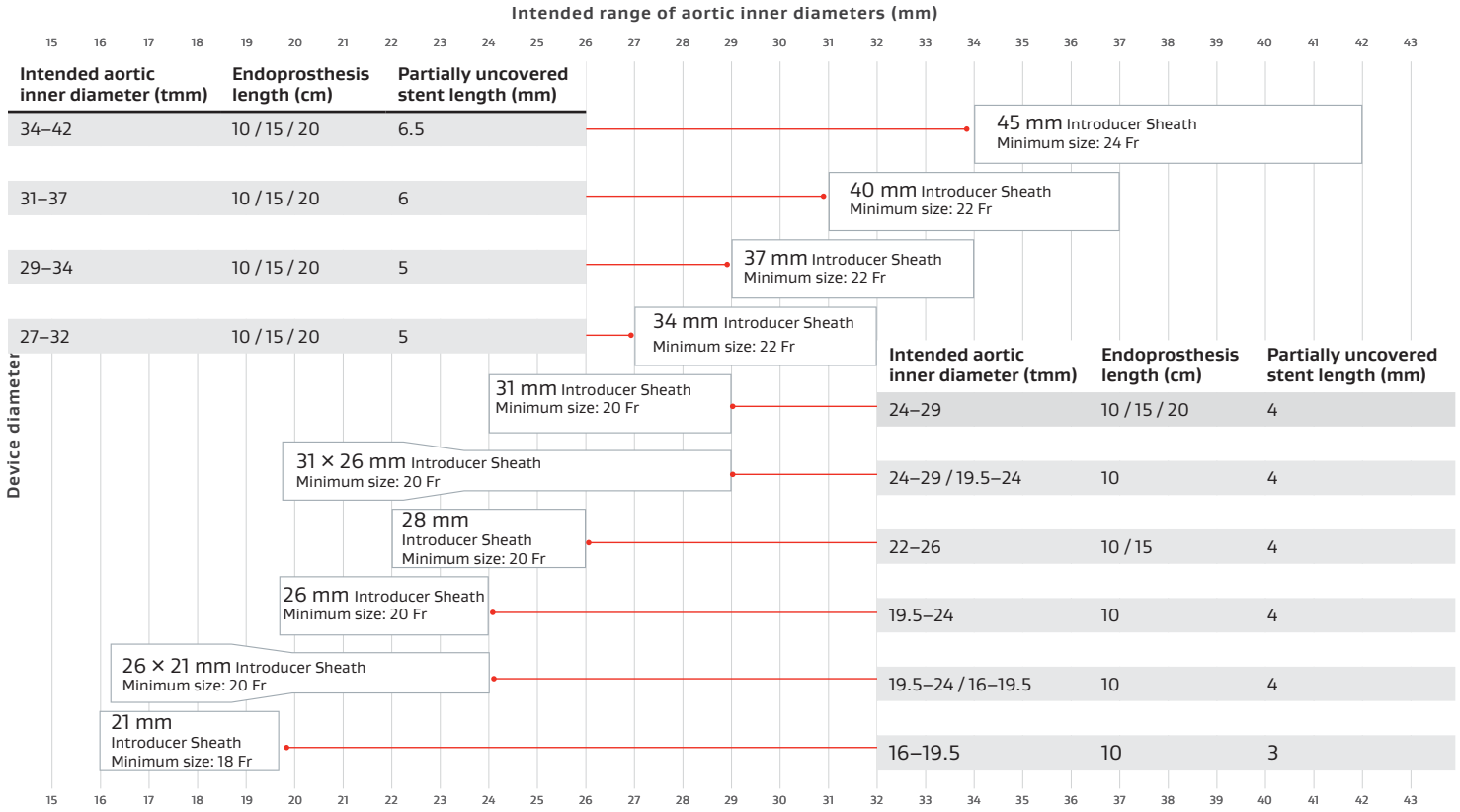
**GORE® TAG®**
Conformable Thoracic Stent Graft
with ACTIVE CONTROL System

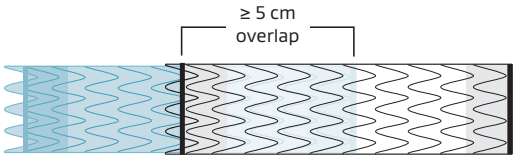
# SIZING GUIDE




| Intended aortic inner diameter (tmm) | Endoprosthesis length (cm) | Partially uncovered stent length (mm) |
|---|---|---|
| 34–42 | 10 / 15 / 20 | 6.5 |
| 31–37 | 10 / 15 / 20 | 6 |
| 29–34 | 10 / 15 / 20 | 5 |
| 27–32 | 10 / 15 / 20 | 5 |
| 24–29 | 10 / 15 / 20 | 4 |
| 24–29 / 19.5–24 | 10 | 4 |
| 22–26 | 10 / 15 | 4 |
| 19.5–24 | 10 | 4 |
| 19.5–24 / 16–19.5 | 10 | 4 |
| 16–19.5 | 10 | 3 |

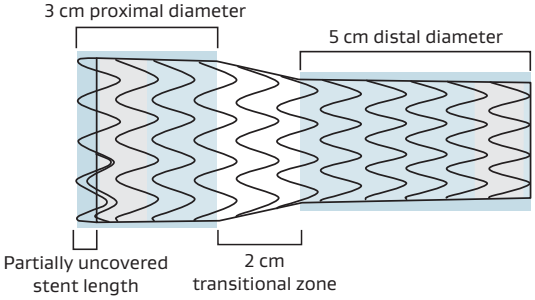
## Using multiple devices

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

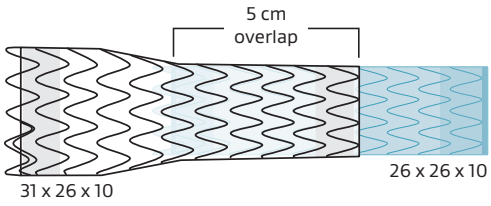


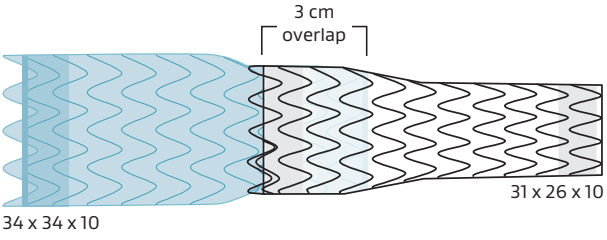

## Tapered devices

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




## Overlapping a tapered device




*Together, improving life*

## 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) |
|---|---|---|---|
| DSF1833 | 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. $R_{X\ Only}$ **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. $R_{X\ Only}$

Products listed may not be available in all markets.



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