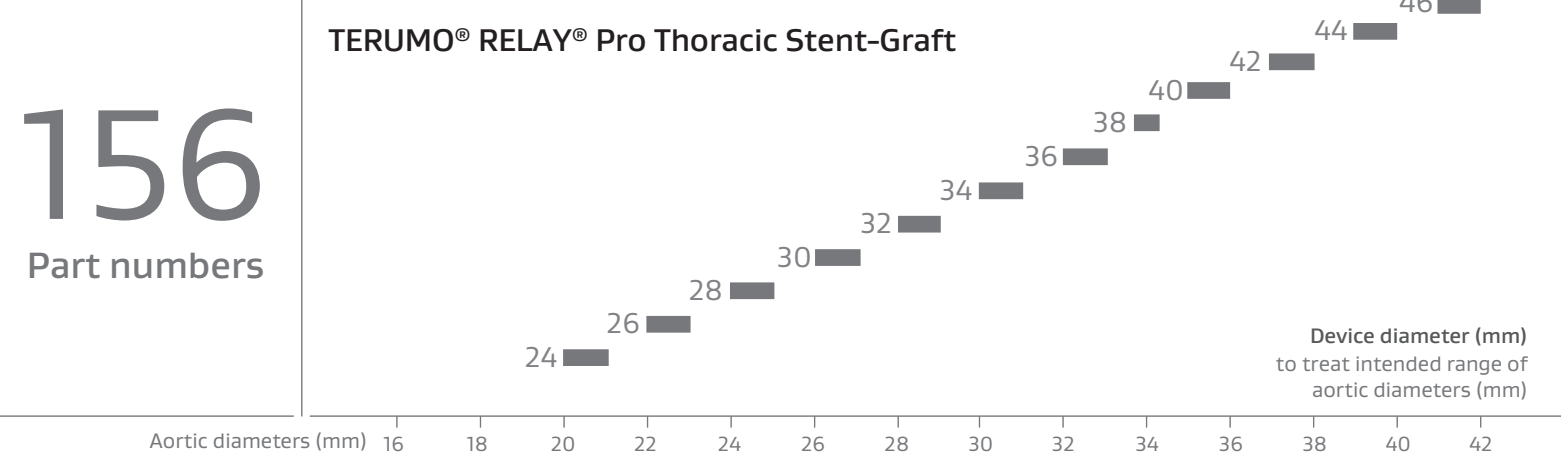
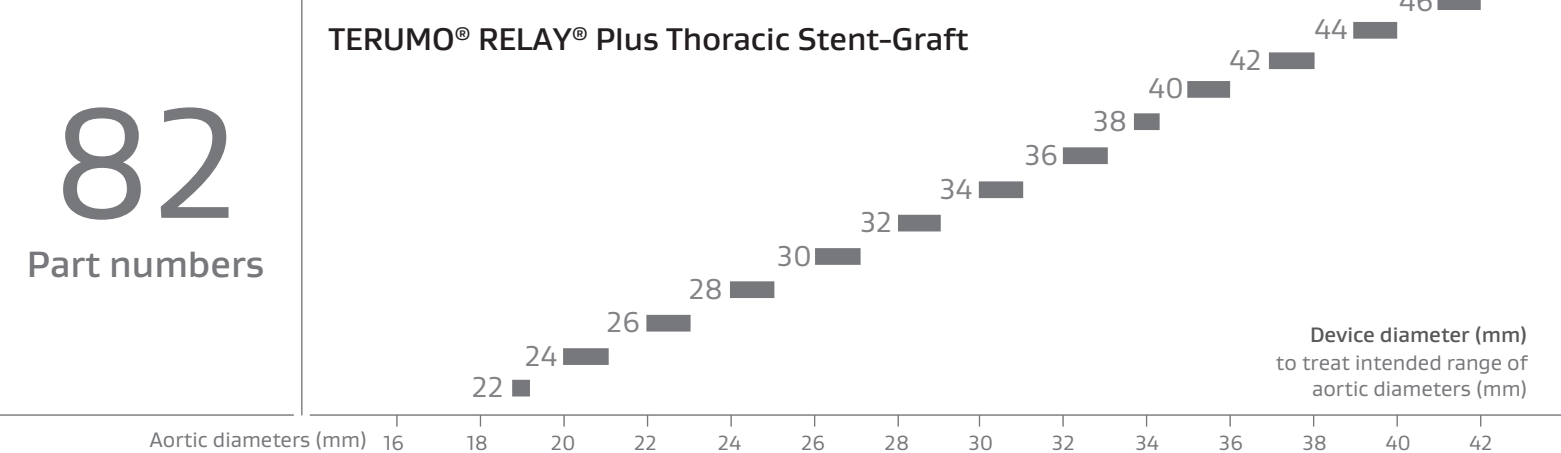
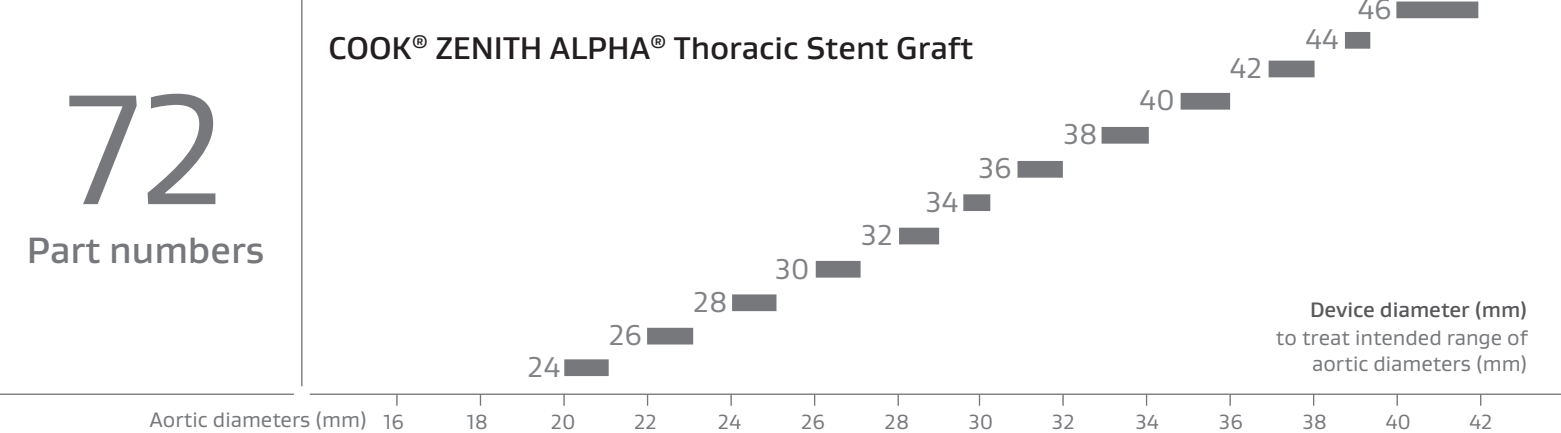
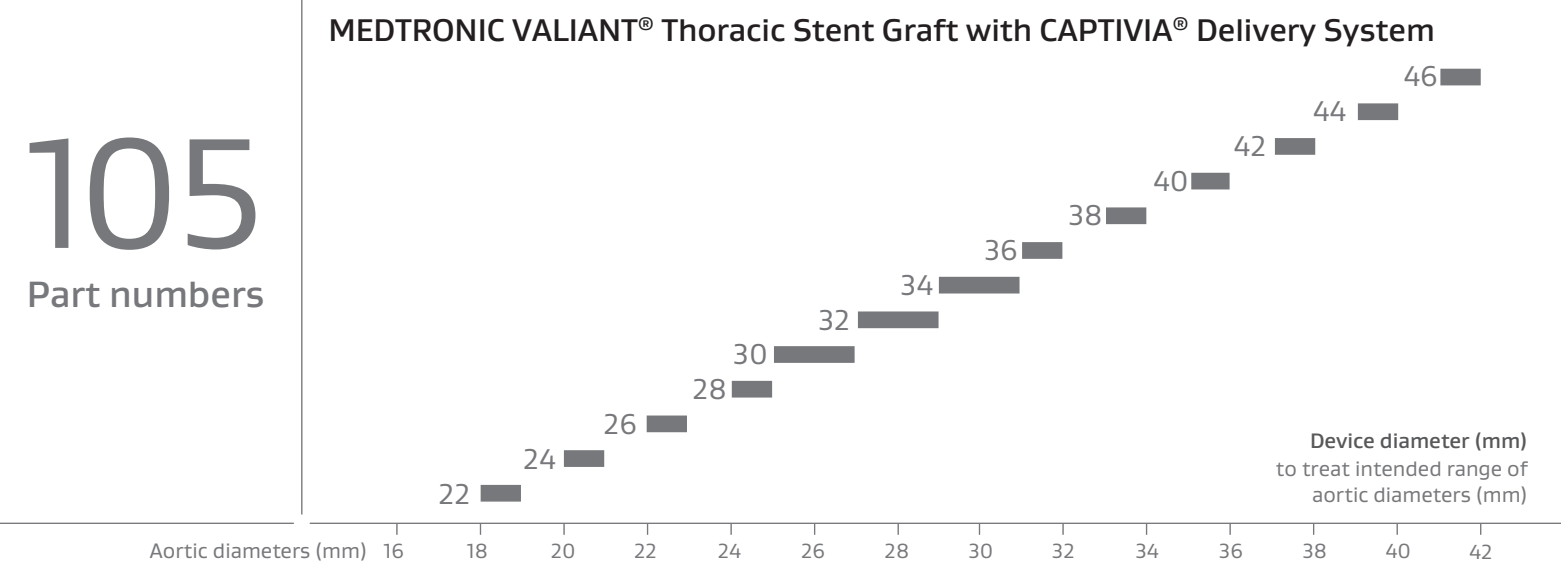
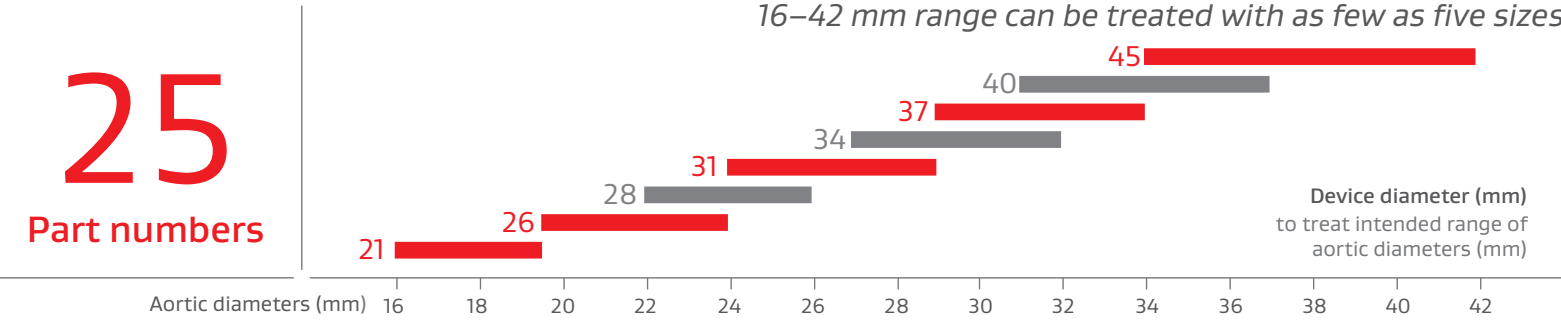


GORE® TAG® Conformable Thoracic Stent Graft with ACTIVE CONTROL System



Note: Device sizing for all devices listed assumes rounding of measured vessels diameter (mm) to nearest whole number within IFU sizing range for aneurysm. For complete information on devices selection and use, please consult the product packaging labels and *Instructions for Use* (IFU). Device sizes and recommendations do not necessarily correlate to clinical outcomes. The material is based on currently available information and may be updated by the manufacturers at any time. Listed products have been cleared or approved by respective regulatory agencies.

 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

Products listed may not be available in all markets.

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