GORE® EXCLUDER®
Conformable AAA Endoprosthesis
with ACTIVE CONTROL System

Trunk-Ipsilateral Leg Endoprosthesis

Catalogue number*	Intended aortic inner diameter (mm)	Aortic endoprosthesis diameter (mm)	Intended iliac inner diameter (mm)	Iliac endoprosthesis diameter (mm)	Overall device length (cm)	Recommended sheath size (Fr)
CXT201212	16–18	20	10–11	12	12	15
CXT201412	16–18	20	12–13.5	14.5	12	15
CXT231412	19–21	23	12–13.5	14.5	12	15
CXT261412	22–23	26	12–13.5	14.5	12	16
CXT281412	24–26	28.5	12–13.5	14.5	12	16
CXT321414	27–29	32	12–13.5	14.5	14	18
CXT361414	30–32	36	12–13.5	14.5	14	18

Aortic Extender Endoprosthesis

Catalogue number*	Intended aortic inner diameter (mm)	Endoprosthesis diameter (mm)	Endoprosthesis length (cm)	Recommended sheath size (Fr)
CXA200005	16–18	20	4.5	15
CXA230005	19–21	23	4.5	15
CXA260005	22–23	26	4.5	15
CXA280005	24–26	28.5	4.5	16
CXA320005	27–29	32	4.5	18
CXA360005	30–32	36	4.5	18

* Currently available commercially.

Together, improving life

Consult Instructions for Use
eifu.goremedical.com

INDICATIONS FOR USE: The GORE® EXCLUDER® Conformable AAA Endoprosthesis is intended to exclude the aneurysm from the blood circulation in patients diagnosed with infrarenal abdominal aortic aneurysm (AAA) disease and who have appropriate anatomy as described below: Adequate iliac/femoral access; Infrarenal aortic neck treatment diameter range of 16–32 mm and a minimum aortic neck length of 10 mm; Proximal aortic neck angulation ≤ 90°; Iliac artery treatment diameter range of 8–25 mm and iliac distal vessel seal zone length of at least 10 mm. **Aortic Extender Endoprosthesis and Iliac Extender Endoprosthesis Components** are intended to be used after deployment of the GORE® EXCLUDER® Conformable AAA Endoprosthesis. These extensions are intended to be used when additional length and/or sealing for aneurysmal exclusion is desired. **CONTRAINDICATIONS:** The GORE® EXCLUDER® Conformable AAA Endoprosthesis is contraindicated in: patients with known sensitivities or allergies to the device materials. All components of the GORE® EXCLUDER® Conformable Endoprosthesis contain expanded polytetrafluoroethylene (ePTFE), fluorinated ethylene propylene (FEP), nitinol (nickel-titanium alloy) and gold. Patients with 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. ℞ Only

Products listed may not be available in all markets.

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