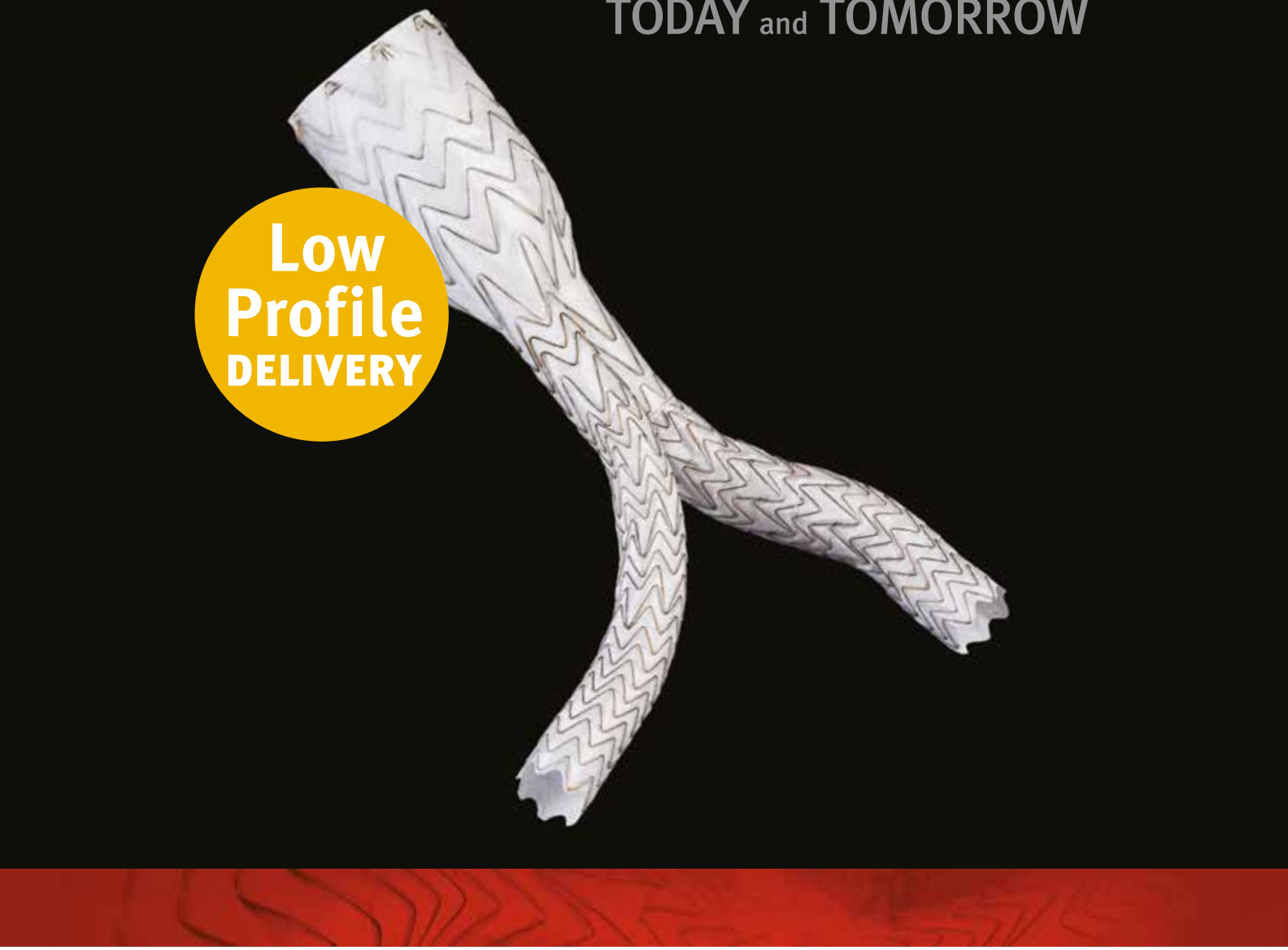
Trusted Performance

TODAY and TOMORROW



GORE EXCLUDER®

AAA ENDOPROSTHESIS

Trusted Design

Time-proven design backed by extensive clinical and commercial data*

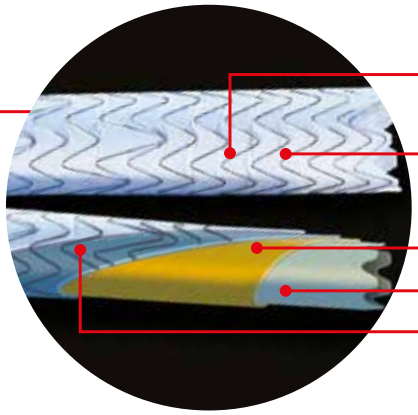
Experience

- Over 20 years worldwide experience
- More than 300,000 devices distributed

Durability

1. **Sutureless construction**
   - Expanded PTFE graft technology on luminal and abluminal surfaces
2. **Advanced sinusoidal stent design**
   - Enhances flexibility and long-term conformability
3. **Proprietary ePTFE film layers**
   - Low permeability with abrasion-resistant properties
   - Conformability in tortuous anatomies
4. **Sealing cuff**
   - Engineered to provide security against endoleaks
5. **Active infrarenal fixation**
   - Anchors for active fixation are engineered to provide migration resistance

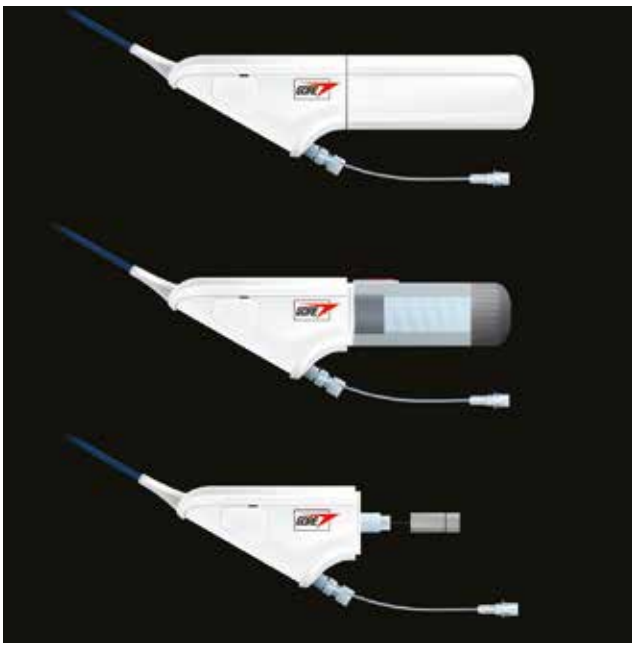
* Data on file



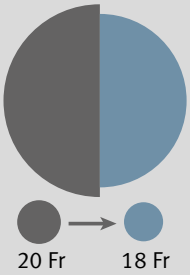
Repositionable Delivery System

- Intuitive nested design
- Simple three-stage deployment
- More opportunities to optimize infrarenal seal

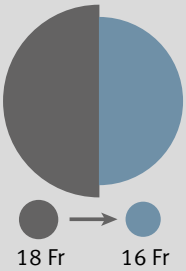
GORE® C3® Delivery System Three-Stage Deployment



Reduced Profile*

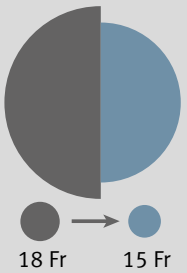


10% Reduction
Trunk-Ipsilateral Leg
31 mm Device

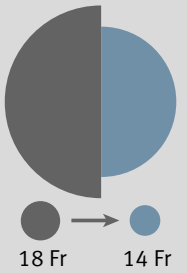


11% Reduction
Trunk-Ipsilateral Leg
23 / 26 mm Device

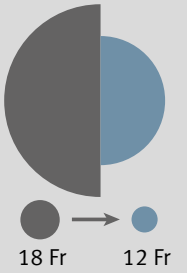
Aortic Extenders
23 / 26 / 28.5 mm



17% Reduction
Contralateral Leg
27 mm Device



22% Reduction
Contralateral Leg
23 mm Device



33% Reduction
Contralateral Leg
16 / 18 / 20 mm Device

* Compared to prior sizes

Expanded low profile delivery options

No Changes to the Implant

- Same proven device
- Same ease of use
- Same deployment methodology

Lower Profile Sleeve Properties

- Our proprietary manufacturing process was applied to increase the sleeve strength
- The sleeve allows the device to be held at a smaller crushed diameter

Trusted Portfolio

16 Fr Low Profile Design

- 23 and 26 mm Trunk-Ipsilateral Legs
- 23, 26, and 28.5 mm Aortic Extenders

35 mm Trunk-Ipsilateral Legs

- 18 Fr low profile design
- Expands aortic neck diameter treatment range to 19–32 mm*
- 36 mm Aortic Extender available

23 and 27 mm Contralateral Legs

- 14–15 Fr low profile design
- Expands iliac diameter treatment range to 8–25 mm

* Based on inner-wall to inner-wall measurements

Trunk-Ipsilateral Leg Endoprosthesis

GORE® C3® Delivery System 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)
RLT231212	19–21	23	10–11	12	12	16
RLT231214	19–21	23	10–11	12	14	16
RLT231216	19–21	23	10–11	12	16	16
RLT231218	19–21	23	10–11	12	18	16
RLT231412	19–21	23	12–13.5	14.5	12	16
RLT231414	19–21	23	12–13.5	14.5	14	16
RLT231416	19–21	23	12–13.5	14.5	16	16
RLT231418	19–21	23	12–13.5	14.5	18	16
RLT261212	22–23	26	10–11	12	12	16
RLT261214	22–23	26	10–11	12	14	16
RLT261216	22–23	26	10–11	12	16	16
RLT261218	22–23	26	10–11	12	18	16
RLT261412	22–23	26	12–13.5	14.5	12	16
RLT261414	22–23	26	12–13.5	14.5	14	16
RLT261416	22–23	26	12–13.5	14.5	16	16
RLT261418	22–23	26	12–13.5	14.5	18	16
RLT281212	24–26	28.5	10–11	12	12	18
RLT281214	24–26	28.5	10–11	12	14	18
RLT281216	24–26	28.5	10–11	12	16	18
RLT281218	24–26	28.5	10–11	12	18	18
RLT281412	24–26	28.5	12–13.5	14.5	12	18
RLT281414	24–26	28.5	12–13.5	14.5	14	18
RLT281416	24–26	28.5	12–13.5	14.5	16	18
RLT281418	24–26	28.5	12–13.5	14.5	18	18
RLT311413	27–29	31	12–13.5	14.5	13	18
RLT311415	27–29	31	12–13.5	14.5	15	18
RLT311417	27–29	31	12–13.5	14.5	17	18
RLT351414	30–32	35	12–13.5	14.5	14	18
RLT351416	30–32	35	12–13.5	14.5	16	18
RLT351418	30–32	35	12–13.5	14.5	18	18

Contralateral Leg Endoprosthesis

Catalogue Number	Intended iliac inner diameter (mm)	Iliac Endoprosthesis Diameter (mm)	Contralateral Leg Length (cm)	Recommended Sheath Size (Fr)
PLC121000	10–11	12	10	12
PLC121200	10–11	12	12	12
PLC121400	10–11	12	14	12
PLC141000	12–13.5	14.5	10	12
PLC141200	12–13.5	14.5	12	12
PLC141400	12–13.5	14.5	14	12
PLC161000*	13.5–14.5	16	9.5	12
PLC161200*	13.5–14.5	16	11.5	12
PLC161400*	13.5–14.5	16	13.5	12
PLC181000*	14.5–16.5	18	9.5	12
PLC181200*	14.5– 16.5	18	11.5	12
PLC181400*	14.5–16.5	18	13.5	12
PLC201000*	16.5–18.5	20	9.5	12
PLC201200*	16.5–18.5	20	11.5	12
PLC201400*	16.5–18.5	20	13.5	12
PLC231000*	18.5–21.5	23	10	14
PLC231200*	18.5–21.5	23	12	14
PLC231400*	18.5–21.5	23	14	14
PLC271000*	21.5–25.0	27	10	15
PLC271200*	21.5–25.0	27	12	15
PLC271400*	21.5–25.0	27	14	15

GORE® DrySeal Sheath with Hydrophilic Coating

Catalogue Number	Sheath Size (Fr)
DSL1228	12
DSL1428	14
DSL1628	16
DSL1828	18

All sheaths are 28 cm in length.

GORE® DrySeal Flex Sheath

Catalogue Number	Sheath Size (Fr)
DSF1233	12
DSF1433	14
DSF1533	15
DSF1633	16
DSF1833	18

All sheaths are 33 cm in length.

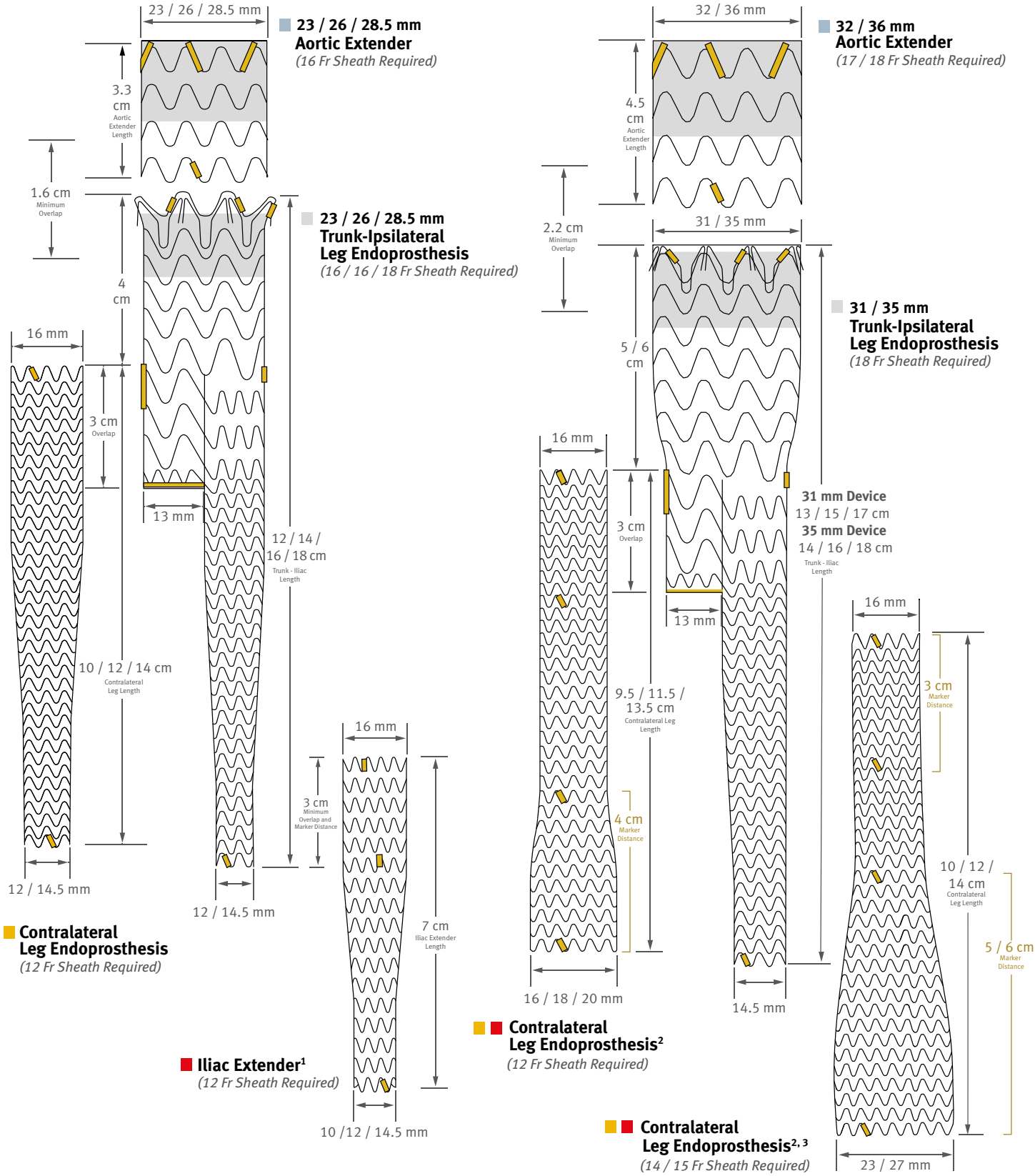
Iliac Extender Endoprosthesis

Catalogue Number	Intended Iliac Inner Diameter (mm)	Endoprosthesis Diameter (mm)	Endoprosthesis Length (cm)	Recommended Sheath Size (Fr)
PLL161007	8–9	10	7	12
PLL161207	10–11	12	7	12
PLL161407	12–13.5	14.5	7	12

Aortic Extender Endoprosthesis

Catalogue Number	Intended Aortic Inner Diameter (mm)	Endoprosthesis Diameter (mm)	Endoprosthesis Length (cm)	Recommended Sheath Size (Fr)
PLA230300	19–21	23	3.3	16
PLA260300	22–23	26	3.3	16
PLA280300	24–26	28.5	3.3	16
PLA320400	27–29	32	4.5	17
PLA360400	30–32	36	4.5	18

* PLEASE NOTE: All large diameter Contralateral Leg Endoprostheses (16, 18, 20, 23, 27 mm) can be used as Iliac Extenders.



Trunk-Ipsilateral Leg
Contralateral Leg
Iliac Extender
Aortic Extender

[1] Traditional sizes of Iliac Extender (10, 12, 14.5 mm) are not compatible with large diameter (16, 18, 20, 23, 27 mm) Contralateral Leg Endoprostheses.
[2] All large diameter Contralateral Leg Endoprostheses (16, 18, 20, 23, 27 mm) can be used as Iliac Extenders.
[3] The 27 mm x 10 cm Contralateral Leg Endoprosthesis does not have a radiopaque marker 6 cm above the distal end.

Service with Integrity

For more than 40 years, we have collaborated with and supported physicians specializing in vascular therapies. We continue that tradition today with our corporate initiatives and field-based associates, who bring value to physicians beyond the innovative design of our products.

Service through Case Support

Pre-case planning and procedural support are focused on patient care.

Service through Physician Education

The Gore MEDICAL MASTERY Series is a commitment to create environments such as Fellows Workshops and Advanced Symposia, where physicians can learn best practices from each other through discussion, case observation, and hands-on work with our cutting-edge simulator technology.

Service through Community Education

In collaboration with physicians, we have created an industry leading program to educate community physicians and patients about AAA risk factors, symptoms, screening and therapies for cardiovascular disease.

GORE
Creative Technologies Worldwide

W. L. Gore & Associates, Inc.
Flagstaff, AZ 86004

+65.67332882 (Asia Pacific)
00800.6334.4673 (Europe)
800.437.8181 (United States)
928.779.2771 (United States)

goremedical.com

INDICATIONS FOR USE: Trunk-Ipsilateral Leg Endoprosthesis and Contralateral Leg Endoprosthesis Components. The GORE® EXCLUDER® 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 19 – 32 mm and a minimum aortic neck length of 15 mm; Proximal aortic neck angulation ≤ 60°; 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.** The Aortic and Iliac Extender Endoprostheses are intended to be used after deployment of the GORE® EXCLUDER® AAA Endoprosthesis. These extensions are intended to be used when additional length and / or sealing for aneurysmal exclusion is desired. **CONTRAINDICATIONS:** The GORE® EXCLUDER® AAA Endoprosthesis is contraindicated in patients with known sensitivities or allergies to the device materials and patients with a systemic infection who may be at increased risk of endovascular graft infection. Refer to *Instructions for Use* at goremedical.com for a complete description of all warnings, precautions and adverse events. ℞ Only

Products listed may not be available in all markets.