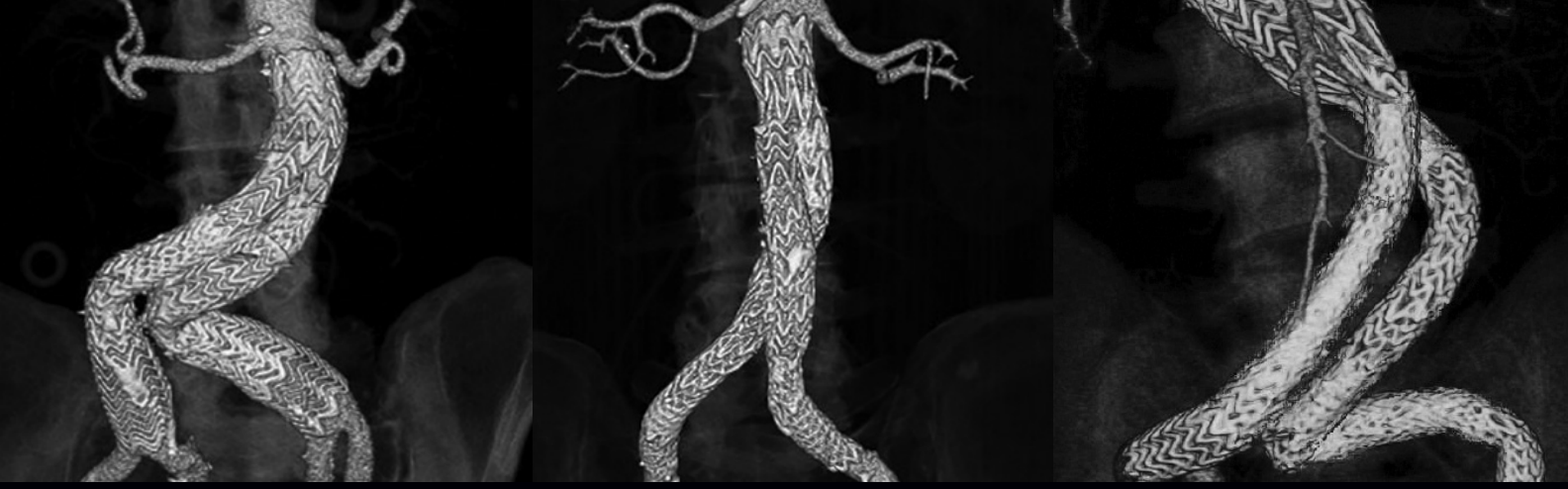
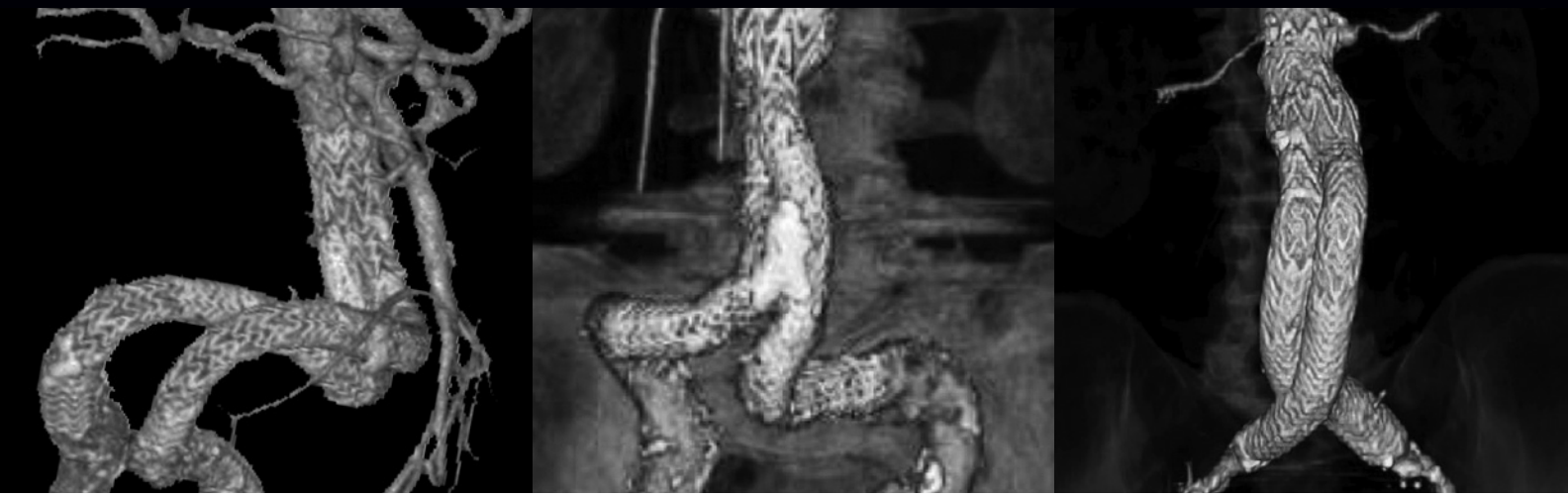


With this much evidence, the choice is clear.



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* Based on company-sponsored trials and registries shown on clinicaltrials.gov for currently available stent grafts.

** GREAT. n = 3,273. To calculate the overall event rates from procedure through end of study period, all subjects who could have had events, regardless of length of follow-up, were included. For outcome data, GREAT only collects site reported serious adverse events.

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GORE EXCLUDER®

AAA ENDOPROSTHESIS

GORE® EXCLUDER® AAA Endoprosthesis

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

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 Iliac 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.

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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.