Durable outcomes. Proven performance.

GORE® EXCLUDER®
AAA Endoprosthesis

GORE® EXCLUDER® Iliac
Branch Endoprosthesis
GORE® EXCLUDER® AAA Endoprosthesis

The most-studied* EVAR stent graft designed for durable outcomes.

The trusted performance of the GORE® EXCLUDER® Device is paired with the intuitive GORE® C3® Delivery System to provide optimal infrarenal seal and reliable results, even in more challenging anatomies.

Proven Performance.
Results from the Global Registry for Endovascular Aortic Treatment (GREAT)

3,273 Patients through 3 years of follow-up**

96.0% Freedom from device-related reintervention

93.4% Freedom from all reintervention

0.0%† Migration

1.4% Type I / III endoleak

0.5% Limb occlusion

GORE® C3® Delivery System

- Repositionable to obtain optimal seal
- Unique ability to reconstrain the proximal end and reposition for ideal placement
- More opportunities to maximize infrarenal seal
GORE® EXCLUDER® Iliac Branch Endoprosthesis

U.S. IDE Clinical Trial now has 2-year follow-up data for all patients from primary enrollment (n = 63).

2-year data

100%  Patency — External iliac artery
95.1%  Patency — Internal iliac artery
93.7%  Freedom from reintervention
0%    Buttock claudication
0%    New onset erectile dysfunction
98.3%  Freedom from CIAA enlargement (> 5 mm)

152 minutes  Procedure time
114 ml      Contrast used
40 minutes  Fluoro time
95.2%  Technical success

We designed this all-in-one system exclusively for use in the iliac arteries. It is the only FDA-approved, off-the-shelf iliac branch solution.

Preservation matters:
Recommended treatment\textsuperscript{2} to sustain quality of life

Performs as promised:
High patency\textsuperscript{3}, conformability, and durability

- Pre-cannulated internal iliac gate and bi-femoral delivery for ease-of-use
- Low profile (16 Fr) delivery for enhanced vessel access and trackability
- Two-staged repositionable deployment for precise placement of the iliac component
The GORE® EXCLUDER® Device family has evolved based on what we have learned from over 20 years of experience in EVAR.

**Worldwide experience**
- More than 300,000 patients treated with GORE® EXCLUDER® AAA Endoprosthesis
- More than 10,000 patients treated with GORE® EXCLUDER® Iliac Branch Endoprosthesis

**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
Physician collaboration and unwavering commitment.

When we join with our partners in the medical community, we work together to advance patient care. With the only FDA-approved iliac branch solution, and durable outcomes with more than 20 years of EVAR experience, the GORE® EXCLUDER® Device family offers solutions physicians trust and patients count on.

Learn more at goremedical.com/aortic

The GORE® EXCLUDER® Iliac Branch Endoprosthesis is intended to be used in conjunction with the GORE® EXCLUDER® AAA Endoprosthesis to isolate the common iliac artery from systemic blood flow and preserve blood flow in the external iliac and internal iliac arteries in patients with a common iliac or aortoiliac aneurysm, who have appropriate anatomy, including: Adequate iliac / femoral access; minimum common iliac diameter of 17 mm at the proximal implantation zone of the IBE; external iliac artery treatment diameter range of 6.5–25 mm and seal zone length of at least 10 mm; internal iliac artery treatment diameter range of 6.5–13.5 mm and seal zone length of at least 10 mm; adequate length from the lowest major renal artery to the internal iliac artery to accommodate the total endoprosthesis length, calculated by adding the minimum lengths of required components, taking into account appropriate overlaps between components. GORE® EXCLUDER® AAA Endoprosthesis Components used in conjunction with GORE® EXCLUDER® Iliac Branch Endoprosthesis: Trunk-Ipsilateral Leg Component. The Trunk-Ipsilateral Leg Extender Endoprostheses can be used after deployment of the GORE® EXCLUDER® AAA Endoprosthesis Components. These extensions are used when additional length and / or sealing for aneurysmal exclusion is desired. CONTRAINDICATIONS: The GORE® EXCLUDER® Iliac Branch Endoprosthesis is contraindicated in: Patients with known sensitivities or allergies to the device materials. All components of the GORE® EXCLUDER® Iliac Branch Endoprosthesis and the GORE® EXCLUDER® AAA Endoprosthesis contain ePTFE, FEP, nitinol (nickel-titanium alloy), and gold. 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. * 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. † One peri-procedural migration reported. Zero migrations reported during follow-up through 3 years. ‡ Defined as successful implantation with lack of endoleaks. § Based on the number of Trunk-Ipsilateral Legs distributed. ‖ Based on the number of Iliac Branch Components distributed.