GORE® EXCLUDER®
Conformable AAA Endoprosthesis
with ACTIVE CONTROL System

BECAUSE EVERY AORTA IS UNIQUE

The latest addition to the GORE® EXCLUDER® Device family offers new levels of control when you need it most.

GET STARTED

Together, improving life

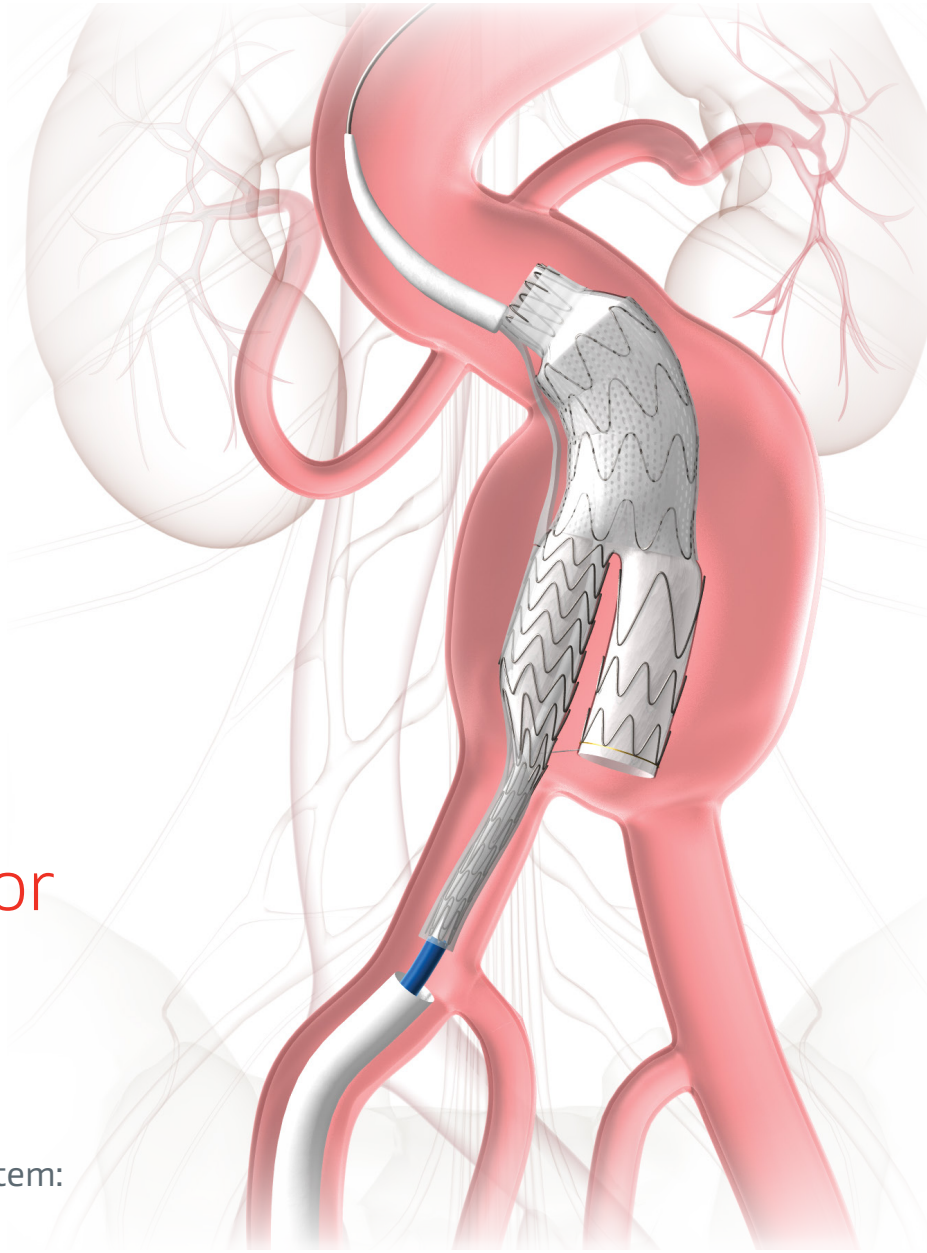
Introducing the GORE® EXCLUDER® Conformable AAA Endoprosthesis with ACTIVE CONTROL System

The conformable stent graft and innovative delivery system are designed to maximize seal.

- Innovation built on a **proven legacy of performance**
- Expands treatment range for small **16–18 mm aortic necks**

The only EVAR device with angulation control

GORE® ACTIVE CONTROL SYSTEM



Controlled conformability for each patient's unique anatomy

The GORE® ACTIVE CONTROL System:

CONFORMABLE STENT GRAFT

- Adapts closely to the anatomy, helping achieve aortic wall apposition.
- Individual stent rows allow for flexibility.

ENHANCED DEVICE POSITIONING

- Ability to reconstrain proximal end for refined positioning.
- Initial trunk deployment at ~70 percent of diameter for ease of use in repositioning.

OPTIONAL ANGULATION CONTROL

- Aids in orthogonal placement to optimize seal within the flow lumen.
- Controlled delivery allows for refinement of angulation at two stages.

Watch the deployment »

PATTERN OF PERFORMANCE

Continuing the pattern of performance

Sub-study outcomes. One-year follow-up.*,†

Zero

- Type I and III endoleaks
- Migrations
- Conversions to open repair
- Ruptures
- Stent fractures
- Limb occlusions

100%

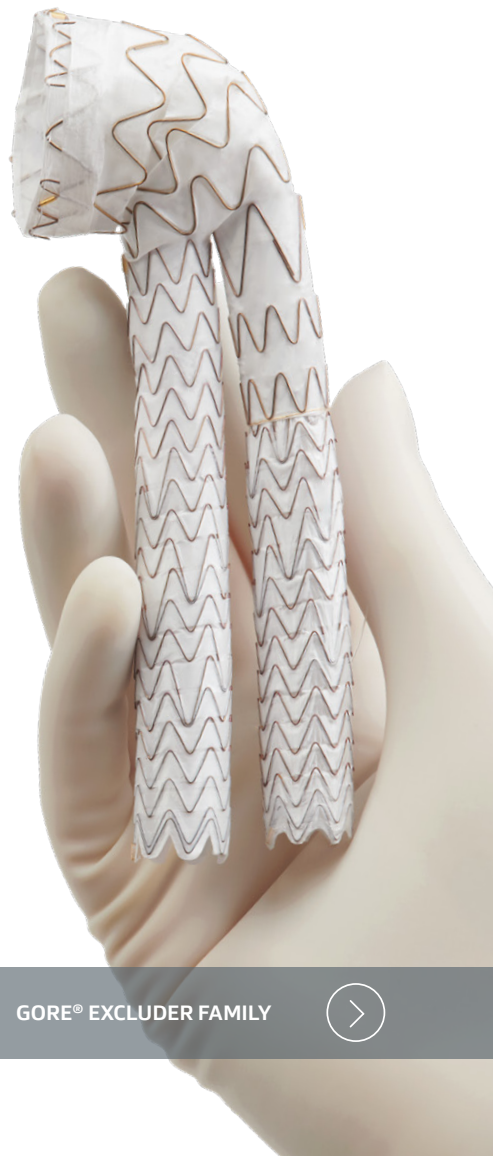
Technical success

Freedom from device-related serious adverse events

Patency

98.6%

Freedom from aneurysm enlargement



* 1-year follow-up. Assessment of the GORE® EXCLUDER® Conformable AAA Endoprosthesis in the Treatment of Abdominal Aortic Aneurysms. NLM Identifier: NCT02489539. Published July 3, 2015. Updated June 16, 2020. Accessed November 17, 2020. https://clinicaltrials.gov/ct2/show/NCT02489539

† For these data points, a minimum of 66 patients were eligible for 1-year outcome analysis, meeting all follow-up requirements that included contrast enhanced CT's. More than 66 patients were included in some data points, which can be confirmed in the *Instructions for Use*.

GORE® EXCLUDER FAMILY

The GORE® EXCLUDER® Conformable AAA Device builds on the proven performance of the GORE® EXCLUDER® Device family

Most EVAR device STUDIED*

450,000+
patients treated†

20+ years
of worldwide experience

The only FDA-approved
iliac branch solution

GORE® EXCLUDER®
Iliac Branch Endoprosthesis

GORE® EXCLUDER®
AAA Endoprosthesis
featuring C3® Delivery System

GORE® EXCLUDER®
Conformable AAA Endoprosthesis
with ACTIVE CONTROL System

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

† Based on the number of trunk-ipsilateral legs distributed for GORE® EXCLUDER® AAA Endoprosthesis.

CONTACT US

The EVAR solutions that set the bar for excellence within aortic treatment

We are continuously innovating to eliminate yesterday's limitations, so you can open new possibilities in patient care.

Contact a Gore representative to learn more.

goremedical.com/EVARangulation

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

INDICATIONS FOR USE IN THE U.S.: The GORE® EXCLUDER® Iliac Branch Endoprosthesis (IBE) is intended to be used 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 is intended to provide proximal seal and fixation for the endovascular repair of the aneurysm. Contralateral Leg Endoprosthesis Component. The Contralateral Leg Endoprosthesis is intended to bridge the GORE® EXCLUDER® Device Trunk-Ipsilateral Component to the GORE® EXCLUDER® Iliac Branch Endoprosthesis following deployment of the GORE® EXCLUDER® Iliac Branch Endoprosthesis. Additionally, the Contralateral Leg Endoprosthesis is intended to be used for distal extension of the Iliac Branch Component in the external iliac artery. The Iliac Branch Component can treat external iliac artery diameters up to 13.5 mm. This ability to extend the Iliac Branch Component distally with any Contralateral Leg Endoprosthesis expands the external iliac artery treatment range up to 25 mm. Aortic Extender Endoprosthesis and Iliac Extender Endoprosthesis Components: The Aortic and Iliac Extender Endoprostheses can be used after deployment of the GORE® EXCLUDER® Iliac Branch Endoprosthesis and GORE® EXCLUDER® AAA Endoprosthesis. 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 eifu.goremedical.com for a complete description of all applicable indications, warnings, precautions and contraindications for the markets where this product is available. ℞ Only

GORE® EXCLUDER® Conformable AAA Endoprosthesis

INDICATIONS FOR USE IN THE U.S.: Trunk-Ipsilateral Leg Endoprosthesis and Contralateral Leg Endoprosthesis Components: 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: Adequate iliac/femoral access, infrarenal aortic neck treatment diameter range of 16–32 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® 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 and patients with a 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

Consult Instructions for Use
eifu.goremedical.com

Products listed may not be available in all markets.

W. L. Gore & Associates, Inc.
goremedical.com

Asia Pacific +65.67332882 **Australia/New Zealand** 1800 680 424 **Europe** 00800 6334 4673
United States Flagstaff, AZ 86004 800 437 8181 928 779 2771