Infrarenal by Choice.

Suprarenal Fixation Devices
- Renal Impairment Risk
- Additional Procedural Steps
- “Top Cap” Deployment Issues
- Suprarenal Angulation Restrictions
- Open Conversion Issues

GORÉ® EXCLUDER®
AAA Endoprosthesis
STAY PUT, STAY PATENT,
STAY INFRArenal.

Outstanding migration and patency data with
0.80% migration and 1.98% renal impairment.

A PubMed® search of all EVAR literature since 2001 generated 7 peer-reviewed publications¹ that report on kidney function or renal artery complications following EVAR and met basic criteria for inclusion in this literature review. The inclusion criteria for these publications was a minimum of 40 patients and follow-up of greater than one year.

This literature review is intended to reflect real-world experience as presented by physicians in the literature, and is not intended as a substitute or comparison to clinical trial data.

¹For the complete list of references, please visit www.goremedical.com/infrarenalbychoice.


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 – 29 mm and a minimum aortic neck length of 15 mm; Proximal aortic neck angulation ≤ 60°; Iliac artery treatment diameter range of 8 – 18.5 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.