

GORE® VIABAHN® VBX Balloon Expandable Endoprosthesis

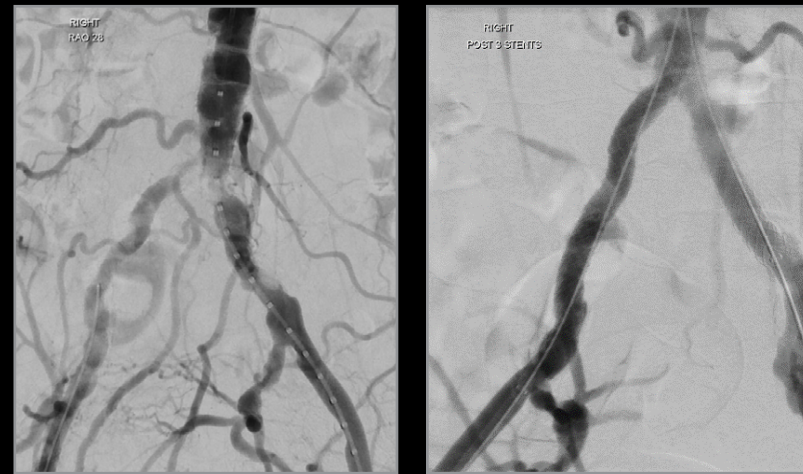


Flexible Strength. Proven Success.



BALLOON EXPANDABLE
ENDOPROSTHESIS

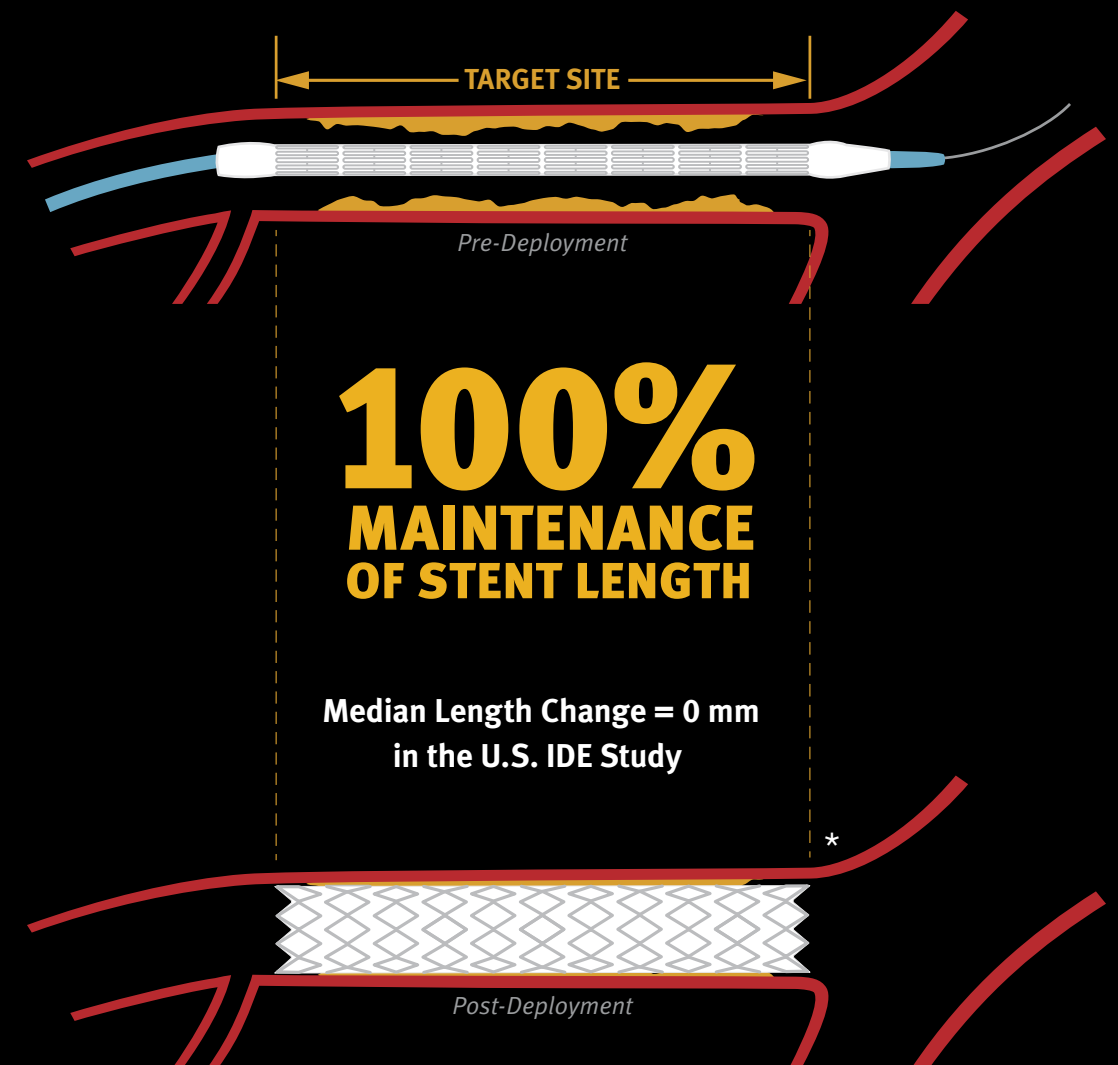
100% RESTORATION OF LUMEN DIAMETER



Before

After

≤ 30% residual stenosis due to high radial strength, even in highly calcified and non-compliant lesions*

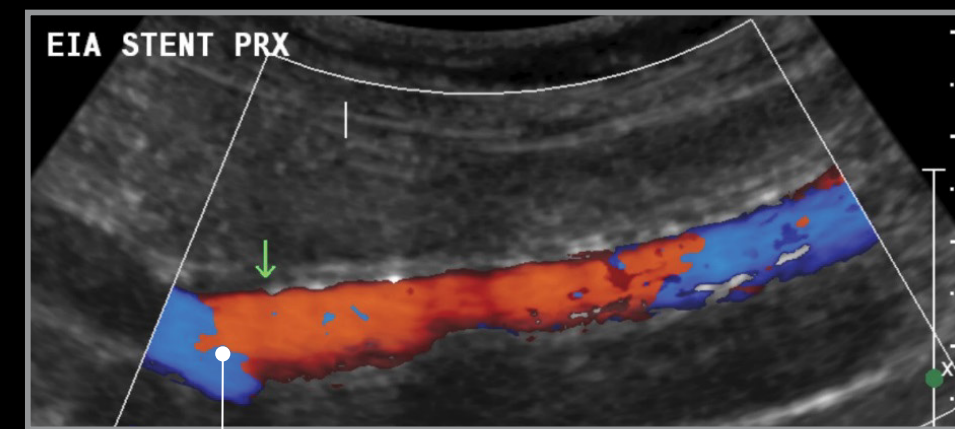


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100% STENT DELIVERY

- 100% delivery to target lesion
- 100% stent retention
- 100% deployment at the target site
- 234 devices delivered
 - 50% bilateral treatment
 - 42% kissing stents
 - 32% TASC II C & D including occlusions
 - 18% contralateral deliveries
 - predilatation not required

96.9% PRIMARY PATENCY



95.3% Primary Patency in
TASC II C & D lesions

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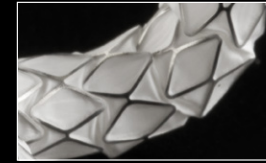


Procedural economic value

GORE® VIABAHN® VBX Balloon Expandable Endoprosthesis (VBX Stent Graft) delivers an estimated savings of \$1,124 / case**

- 79 mm length reduces the number of devices needed in certain instances
- Not required to pre-dilate with the VBX Stent Graft
- Reliable delivery with no device dislodgements
- Accurate delivery helps avoid stent foreshortening and elongation

Advanced technology and unique design



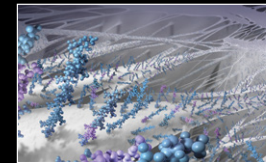
Stainless steel independent rings enhance flexibility

- Minimizes foreshortening
- Provides high radial strength



Semi-compliant covered balloon

- Enables diameter customization
- Improves device retention and trackability in torturous anatomies



CBAS Heparin Surface for lasting thromboresistance



Expanded range of diameters and lengths



Proven leader with 20 years of peripheral stent graft clinical experience

- Employs the covered stent technology of GORE® VIABAHN® Endoprosthesis

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See flexible strength in action:
goremedical.com/vbx-expands



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* Data on file. Based on nine month results from the Gore VBX FLEX IDE Clinical Study (N=134 subjects) and follow-ups conducted for US FDA approval.
** Data on file.

INTENDED USE / INDICATIONS: The GORE® VIABAHN® VBX Balloon Expandable Endoprosthesis is indicated for the treatment of *de novo* or restenotic lesions found in iliac arteries with reference vessel diameters ranging from 5 mm – 13 mm and lesion lengths up to 110 mm, including lesions at the aortic bifurcation. **CONTRAINDICATIONS:** Do not use the GORE® VIABAHN® VBX Balloon Expandable Endoprosthesis in patients with known hypersensitivity to heparin, including those patients who have had a previous incident of Heparin-Induced Thrombocytopenia (HIT) type II. Refer to *Instructions for Use* at goremedical.com for a complete description of all contraindications, warnings, precautions and adverse events. Rx Only

Products listed may not be available in all markets.

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