GORE® VIABAHN® VBX
Balloon Expandable
Endoprosthesis

FLEXIBLE STRENGTH. PROVEN SUCCESS.

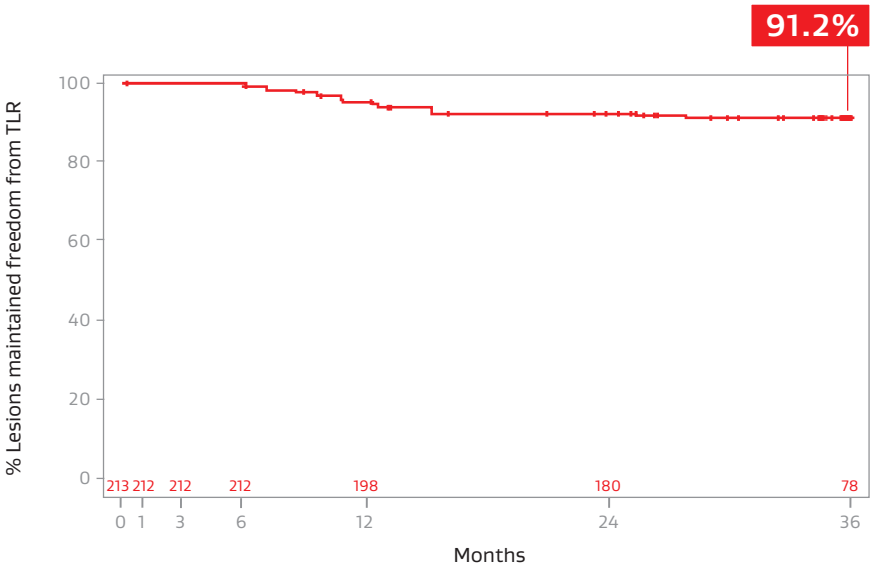
Proven procedural success and durable clinical outcomes through 3 years

Proven procedural success:

- 100% restoration of lumen diameter[1]
- 100% delivery to target lesion with no device dislodgement[1]
- 96.9% primary patency at nine months[1]

Sustained clinical effectiveness through 3 years:

- 91.2% freedom from target lesion revascularization (fTLR)[2]



Kaplan-Meier graph of fTLR per lesion with number of lesions at risk

The Gore VBX FLEX Clinical Study is a prospective, multicenter, single-arm study of 134 patients with complex aortoiliac occlusive disease (32.1% TASC II C and D, 42.5% kissing stent)

Together, improving life

DURABLE PATIENT BENEFIT VERSUS BASELINE THROUGH 3 YEARS[2]

92% **of patients with improvement in Rutherford category[2]**

.17 **improvement in mean resting ankle-brachial index (ABI) (*P* < .001, .93 mean ABI)[*,2]**

2-3x improvement in median WIQ measures[†]

	Pre-procedure	9 months	2 years	3 years
Walking distance	8 (N = 134)	22 (N = 114)	22 (N = 104)	22 (N = 90)
Walking speed	3 (N = 134)	11 (N = 114)	10 (N = 104)	10 (N = 90)
Stair climbing	4 (N = 127)	9 (N = 114)	9 (N = 102)	9 (N = 87)

* (*P* < .001) Statistically significant change from pre-procedure.

† Data on file 2020; W. L. Gore & Associates, Inc.; Flagstaff, AZ.

References

1. Bismuth J, Gray BH, Holden A, Metzger C, Panneton J; VBX FLEX Study Investigators. Pivotal study of a next-generation balloon-expandable stent-graft for treatment of iliac occlusive disease. *Journal of Endovascular Therapy* 2017;24(5):629-637. http://journals.sagepub.com/doi/full/10.117VB7/1526602817720463.

2. Panneton JM, Bismuth J, Gray BH, Holden A. Three-year follow-up of patients with iliac occlusive disease treated with the Viabahn Balloon-Expandable Endoprosthesis. *Journal of Endovascular Therapy*. In press. https://journals.sagepub.com/doi/10.1177/1526602820920569

INDICATIONS FOR USE IN THE U.S., AUSTRALIA, NEW ZEALAND, CANADA AND LATIN AMERICA: 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 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.

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