

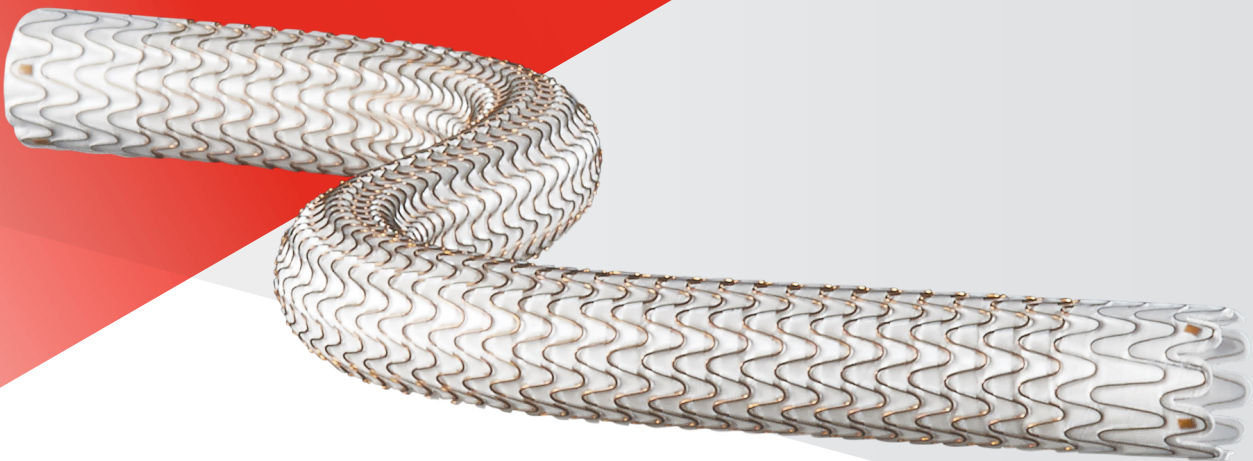


**GORE® VIABAHN®**

Endoprosthesis  
with PROPATEN Bioactive Surface\*

PROVEN PATENCY.†  
DEMONSTRATED DURABILITY.†

Gore Japan Post-Market Clinical Study



\* As used by Gore, PROPATEN Bioactive Surface refers to Gore's proprietary CBAS® Heparin Surface.

† GORE® VIABAHN® Endoprosthesis. W. L. Gore & Associates website. Accessed October 24, 2023.  
<https://www.goremedical.com/VIABAHN/references>.

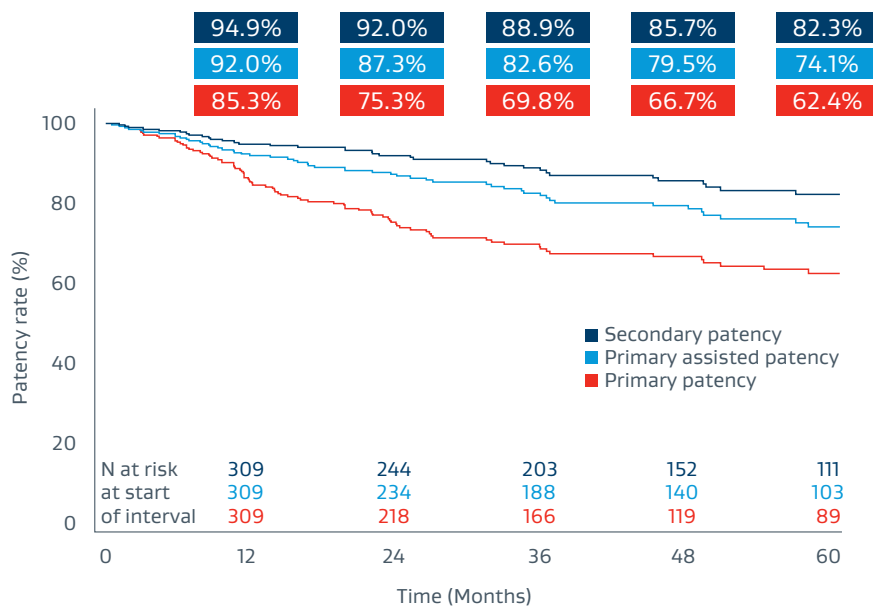
# Gore Japan Post-Market Clinical Study results: Durable clinical outcomes through 5 years

Complex, real-world patient population with challenging superficial femoral artery (SFA) disease<sup>1</sup>:

- 24 cm average lesion length
- 70% chronic total occlusions (CTO)
- 27% critical limb-threatening ischemia (CLTI)
- 48% TASC II D lesions

## Proven patency

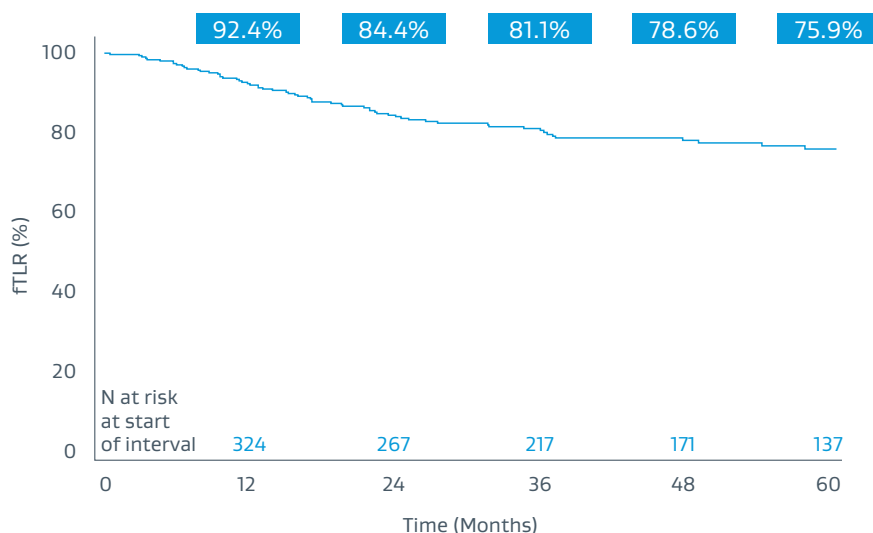
85% primary patency at 1 year, 62% at 5 years<sup>2</sup>



Multivariate analyses did not reveal differences in primary patency for risk factors including lesion length, TASC II class, calcification or CLTI.<sup>2</sup>

## Demonstrated durability

92.4% freedom from target lesion revascularization (fTLR) at 1 year, 75.9% at 5 years<sup>2</sup>



No acute limb ischemia or stent fractures through 5 years.<sup>2</sup>

\* Weighted average lesion length. † One-year weighted average primary patency. ‡ CTO percentage defined as percentage of TASC II D.

Proven patency at 1 year in complex SFA lesions across 7 multicenter, prospective, randomized or single-arm studies<sup>2-8</sup>

1,089

lesions studied

71%

chronic total occlusions (CTO)

23 cm

average lesion length\*

80%

average primary patency†

Trial name	Number of lesions	Mean lesion length (cm)	CTOs (%)	1-year primary patency (%)	1-year secondary patency (%)
SuperB Study <sup>3</sup>	63	23	75 <sup>‡</sup>	65	86
Gore VIPER Clinical Study <sup>4</sup>	119	19	56	73	92
VIASTAR Trial <sup>5</sup>	66	19	79	78	90
25 cm Trial <sup>6</sup>	71	27	93	67	97
Gore Japan IDE Clinical Study <sup>7</sup>	103	22	66	88	98
<b>Gore Japan Post-Market Clinical Study<sup>2</sup></b>	<b>324</b>	<b>24</b>	<b>70</b>	<b>85</b>	<b>95</b>
VANQUISH Study <sup>8</sup>	343	25	71	80	N/A
<b>Combined results (Weighted average, as appropriate)</b>	<b>1,089</b>	<b>23</b>	<b>71</b>	<b>80</b>	<b>94</b>

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8. Iida O, Takahara M, Soga Y, *et al.*; VANQUISH Investigators. One-year outcomes of heparin-bonded stent-graft therapy for real-world femoropopliteal lesions and the association of patency with the prothrombotic state based on the prospective, observational, multicenter Viabahn Stent-Graft Placement for Femoropopliteal Diseases Requiring Endovascular Therapy (VANQUISH) Study. *Journal of Endovascular Therapy* 2021;28(1):123-131.



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