

GORE® VIABAHN®

Endoprosthesis with PROPATEN Bioactive Surface^a

GORE REVISE CLINICAL STUDY

Together, improving life

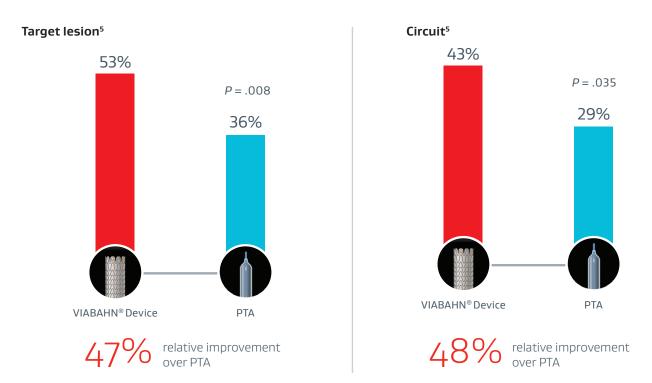
Gore REVISE Clinical Study

Multicenter, prospective, randomized evaluation of the VIABAHN® Device at the venous anastomosis of arteriovenous (AV) grafts versus percutaneous transluminal angioplasty (PTA).

The REVISE Study is the only study to include an endoprosthesis across the elbow and the only randomized control trial to include thrombosed grafts.¹⁻⁴

Results and key takeaways: Patency

The VIABAHN® Device had nearly 50% higher primary patency at 6 months than PTA alone.⁵



Increased primary patency of both the target lesion and the circuit by ~ 50% when compared to PTA at 6 months.⁵

The VIABAHN® Device demonstrated secondary patency across the elbow at 2 years of 83% with 0 reported fractures.6



Results and key takeaways: Thrombosed grafts

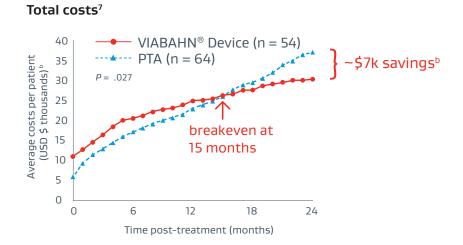
Thrombosed graft lesions treated with the VIABAHN® Device required fewer reinterventions versus PTA at 2 years.⁷



Reinterventions⁷ VIABAHN® Device (n = 54) PTA (n = 64) P = .022 40% Time post-treatment (months)

Fewer reinterventions drove lower costs versus PTA in the US⁷





Compared to total two-year treatment costs with PTA only (\$37,206), treatment with the VIABAHN® Device (\$30,329) becomes more cost-effective at 15 months post-placement, generating \$6,877 in total cost savings per patient.^{b,7}

^b Costs are reflected in 2015 USD

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