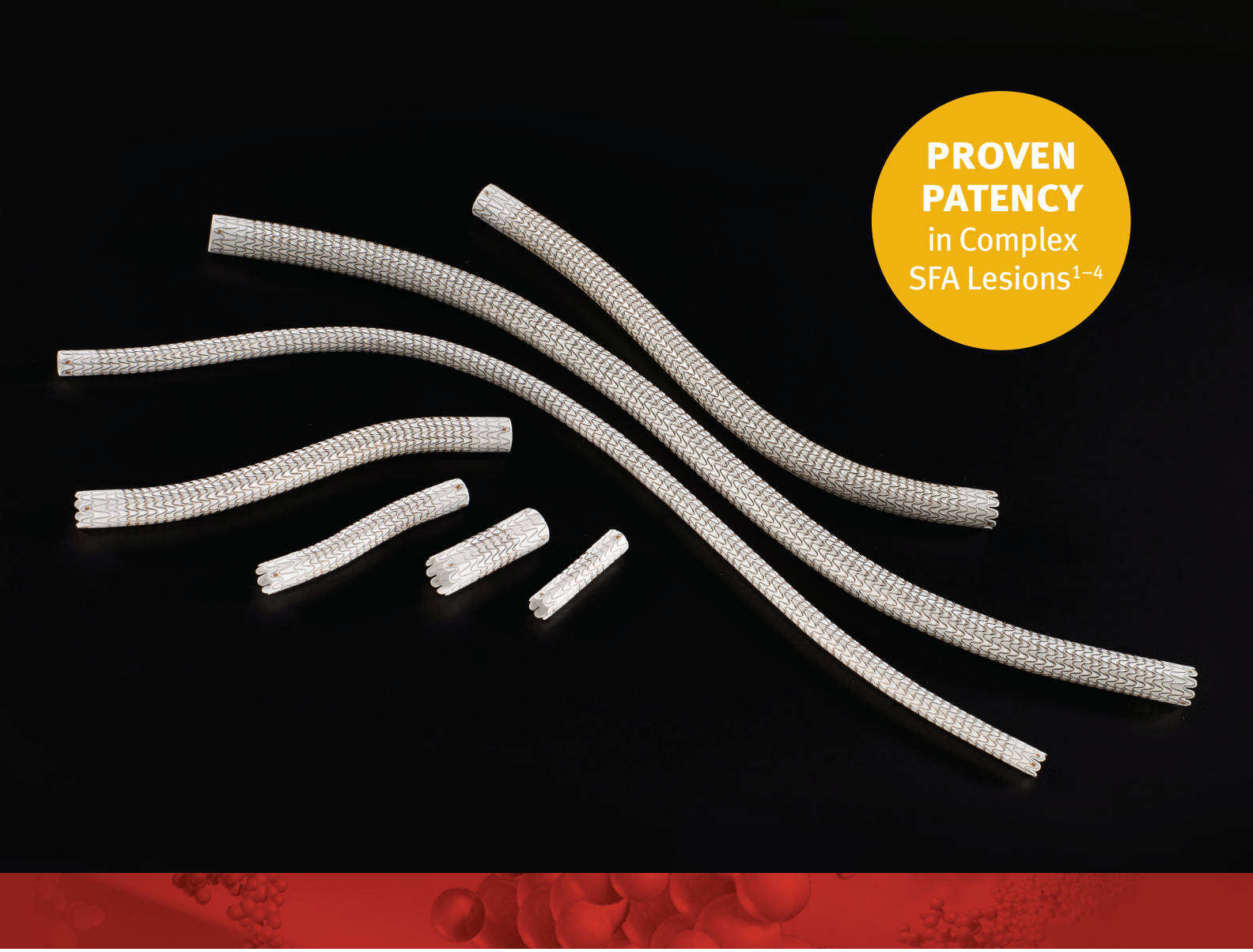

CONFIGURATIONS

GORE

VIABAHN® ENDOPROSTHESIS

PROPATEN BIOACTIVE SURFACE

Catalogue Number	Endoprosthesis Labeled Diameter* (mm)	Endoprosthesis Length* (cm)	Catheter Length** (cm)	Recommended Vessel Diameter† (mm)	Device Profile (Fr)	Recommended Balloon Diameter for Device Touch-up§ (mm)	Guidewire Compatibility
PAJR050202B	5	2.5	120	4.0–4.7	6	5	0.018" or 0.014"
PAJR050502B	5	5.0	120	4.0–4.7	6	5	0.018" or 0.014"
PAJR051002B	5	10.0	120	4.0–4.7	6	5	0.018" or 0.014"
PAJR051502B	5	15.0	120	4.0–4.7	6	5	0.018" or 0.014"
PAJR052502B	5	25.0	120	4.0–4.7	6	5	0.018" or 0.014"
PAJR060202B	6	2.5	120	4.8–5.5	6	6	0.018" or 0.014"
PAJR060502B	6	5.0	120	4.8–5.5	6	6	0.018" or 0.014"
PAJR061002B	6	10.0	120	4.8–5.5	6	6	0.018" or 0.014"
PAJR061502B	6	15.0	120	4.8–5.5	6	6	0.018" or 0.014"
PAJR062502B	6	25.0	120	4.8–5.5	6	6	0.018" or 0.014"
PAJR070202B	7	2.5	120	5.6–6.5	7	7	0.018" or 0.014"
PAJR070502B	7	5.0	120	5.6–6.5	7	7	0.018" or 0.014"
PAJR071002B	7	10.0	120	5.6–6.5	7	7	0.018" or 0.014"
PAJR071502B	7	15.0	120	5.6–6.5	7	7	0.018" or 0.014"
PAJR072502B	7	25.0	120	5.6–6.5	7	7	0.018" or 0.014"
PAJR080202B	8	2.5	120	6.6–7.5	7	8	0.018" or 0.014"
PAJR080502B	8	5.0	120	6.6–7.5	7	8	0.018" or 0.014"
PAJR081002B	8	10.0	120	6.6–7.5	7	8	0.018" or 0.014"
PAJR081502B	8	15.0	120	6.6–7.5	7	8	0.018" or 0.014"
PAJR082502B	8	25.0	120	6.6–7.5	7	8	0.018" or 0.014"
PAH090502B	9	5.0	120	7.6–8.5	9	9	0.035"
PAH091002B	9	10.0	120	7.6–8.5	9	9	0.035"
PAH091502B	9	15.0	120	7.6–8.5	9	9	0.035"
PAH100502B	10	5.0	120	8.6–9.5	11‡	10	0.035"
PAH101002B	10	10.0	120	8.6–9.5	11‡	10	0.035"
PAH101502B	10	15.0	120	8.6–9.5	11‡	10	0.035"
PAH110502B	11	5.0	120	9.6–10.5	11	12	0.035"
PAH111002B	11	10.0	120	9.6–10.5	11	12	0.035"
PAH130502B	13	5.0	120	10.6–12.0	12	14	0.035"
PAH131002B	13	10.0	120	10.6–12.0	12	14	0.035"

Catalogue numbers starting with PAJR indicate low profile items.

* Labeled device diameters and lengths are nominal.

** Ensure the guidewire is the appropriate size (see *Instructions for Use*) and has a length at least twice that of the delivery catheter.

† Recommended endoprosthesis compression within the vessel is approximately 5–20%.

‡ The 10 mm diameter device is compatible with the following 10 Fr introducer sheaths: Cordis AVANTI® Sheath Introducer, Boston Scientific SUPER SHEATH Introducer Sheath, B. Braun INTRADYN Tear-Away Introducer Sheath.

§ For the 11 and 13 mm diameter devices, balloon inflation pressure should not exceed 8 atm.

1. Saxon RR, Chervu A, Jones PA, *et al.* Heparin-bonded, expanded polytetrafluoroethylene-lined stent graft in the treatment of femoropopliteal artery disease: 1-year results of the VIPER (Viabahn Endoprosthesis with Heparin Bioactive Surface in the Treatment of Superficial Femoral Artery Obstructive Disease) Trial. *Journal of Vascular & Interventional Radiology* 2013;24(2):165-173.
2. Lammer J, Zeller T, Hausegger KA, *et al.* Heparin-bonded covered stents versus bare-metal stents for complex femoropopliteal artery lesions: the randomized VIASTAR trial (Viabahn endoprosthesis with PROPATEN bioactive surface [VIA] versus bare nitinol stent in the treatment of long lesions in superficial femoral artery occlusive disease). *Journal of the American College of Cardiology* 2013;62(15):1320-1327.
3. Zeller T, Peeters P, Bosiers M, *et al.* Heparin-bonded stent-graft for the treatment of TASC II C and D femoropopliteal lesions: the Viabahn-25 cm Trial. *Journal of Endovascular Therapy* 2014;21(6):765-774.
4. Ohki T, Kichikawa K, Yokoi H, *et al.* Outcomes of the Japanese multicenter Viabahn trial of endovascular stent grafting for superficial femoral artery lesions. *Journal of Vascular Surgery* 2017;66(1):130-142.e1.

This piece is intended for distribution in Canada and Australia / New Zealand.

Consult Instructions for Use

Products listed may not be available in all markets.

GORE
Creative Technologies Worldwide

W. L. Gore & Associates, Inc.
Flagstaff, AZ 86004

+65.67332882 (Asia Pacific)
00800.6334.4673 (Europe)
800.437.8181 (United States)
928.779.2771 (United States)

goremedical.com