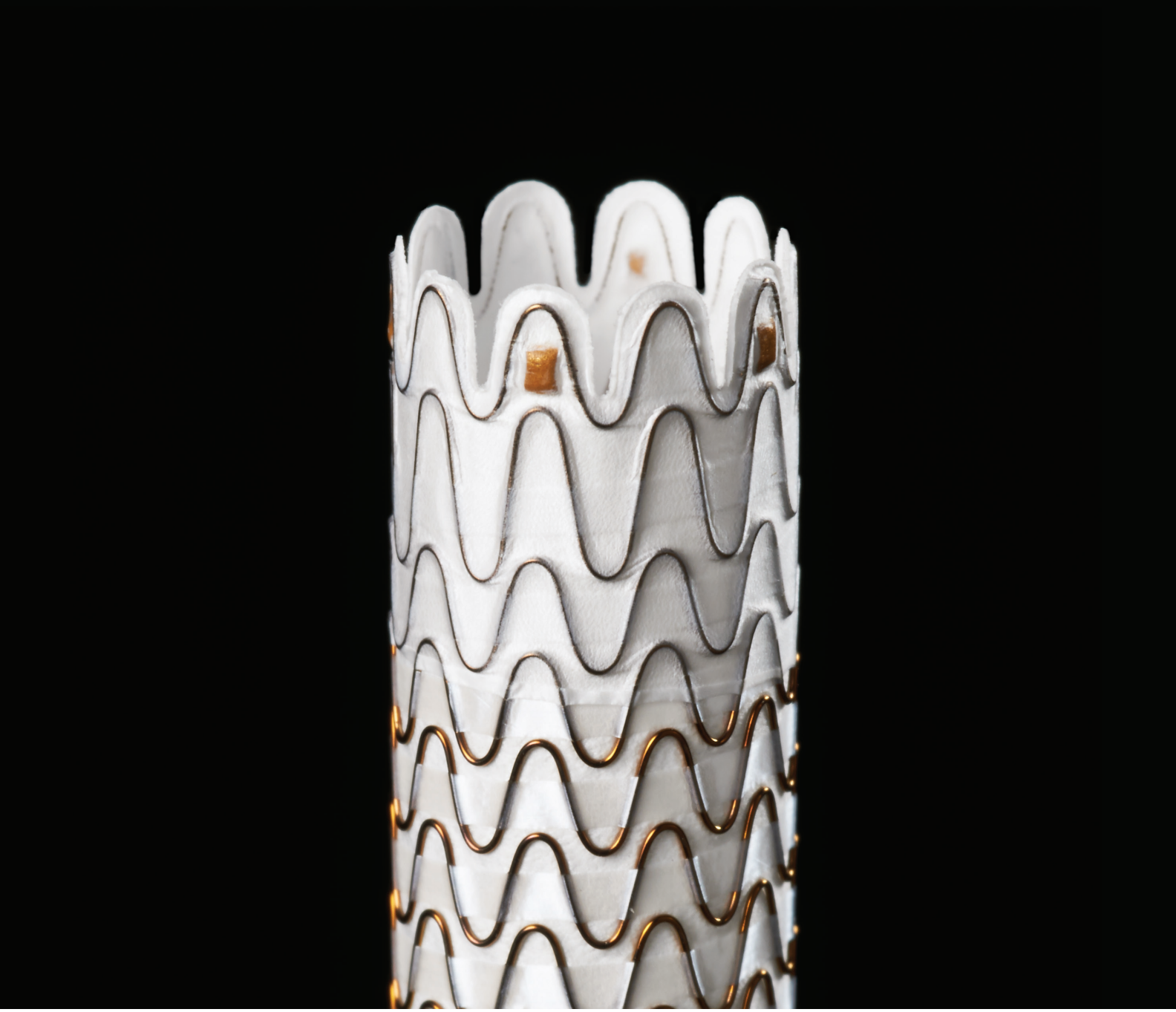

GORE® VIABAHN® Endoprosthesis with Heparin Bioactive Surface*

CONFIGURATIONS

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

SIZING TABLE

.014" / .018" Guidewire compatibility (with radiopaque markers)

Device sizing		Introducer sheath (Fr)			
Endoprosthesis labeled diameter* (mm)	Recommended vessel diameter† (mm)	5 cm Device length*	10 cm Device length*	15 cm Device length*	Recommended balloon diameter for device touch-up‡ (mm)
5	4.0–4.7	6	6	6	5
6	4.8–5.5	6	6	6	6
7	5.6–6.5	7	7	7	7
8	6.6–7.5	7	7	7	8

.035" Guidewire compatibility (with radiopaque markers)

Device sizing		Introducer sheath (Fr)			
Endoprosthesis labeled diameter* (mm)	Recommended vessel diameter† (mm)	5 cm Device length*	10 cm Device length*	15 cm Device length*	Recommended balloon diameter for device touch-up‡ (mm)
9	7.6–8.5	8	8	8	9
10	8.6–9.5	8	8	8	10
11	9.6–10.5	10	10	–	12
13	10.6–12.0	10§	10§	–	14

CATALOGUE LISTINGS

.014" / .018" Guidewire

Catalogue number	Endoprosthesis labeled diameter* (mm)	Endoprosthesis length* (cm)	Catheter length‖ (cm)	Recommended vessel diameter† (mm)	Device profile (Fr)
VBJR050502W	5	5	120	4.0–4.7	6
VBJR051002W	5	10	120	4.0–4.7	6
VBJR051502W	5	15	120	4.0–4.7	6
VBJR060502W	6	5	120	4.8–5.5	6
VBJR061002W	6	10	120	4.8–5.5	6
VBJR061502W	6	15	120	4.8–5.5	6
VBJR070502W	7	5	120	5.6–6.5	7
VBJR071002W	7	10	120	5.6–6.5	7
VBJR071502W	7	15	120	5.6–6.5	7
VBJR080502W	8	5	120	6.6–7.5	7
VBJR081002W	8	10	120	6.6–7.5	7
VBJR081502W	8	15	120	6.6–7.5	7

CATALOGUE LISTINGS

.035" Guidewire

Catalogue number	Endoprosthesis labeled diameter* (mm)	Endoprosthesis length* (cm)	Catheter length ‖ (cm)	Recommended vessel diameter † (mm)	Device profile (Fr)
VBHR090502W	9	5	120	7.6–8.5	8
VBHR091002W	9	10	120	7.6–8.5	8
VBHR091502W	9	15	120	7.6–8.5	8
VBHR100502W	10	5	120	8.6–9.5	8
VBHR101002W	10	10	120	8.6–9.5	8
VBHR101502W	10	15	120	8.6–9.5	8
VBHR110502W	11	5	120	9.6–10.5	10
VBHR111002W	11	10	120	9.6–10.5	10
VBHR130502W	13	5	120	10.6–12.0	10 §
VBHR131002W	13	10	120	10.6–12.0	10 §

* Labeled device diameters and lengths are nominal.
† Recommended endoprosthesis compression within the vessel is approximately 5–20%.
‡ For the 11 and 13 mm diameter devices, balloon inflation pressure should not exceed 8 atm.
§ The 13 mm diameter device is not compatible with the 10 Fr COOK® FLEXOR® CHECK-FLO® Introducers.
‖ Ensure the guidewire is the appropriate size (see *Instructions for Use*) and has a length at least twice that of the delivery catheter.

Consult Instructions for Use
eifu.goremedical.com

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

Products listed may not be available in all markets.

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

+65 67332882 (Asia Pacific)
1800 680 424 (Australia / New Zealand)
00800 6334 4673 (Europe)
800 437 8181 (United States)
928 779 2771 (United States)

goremedical.com