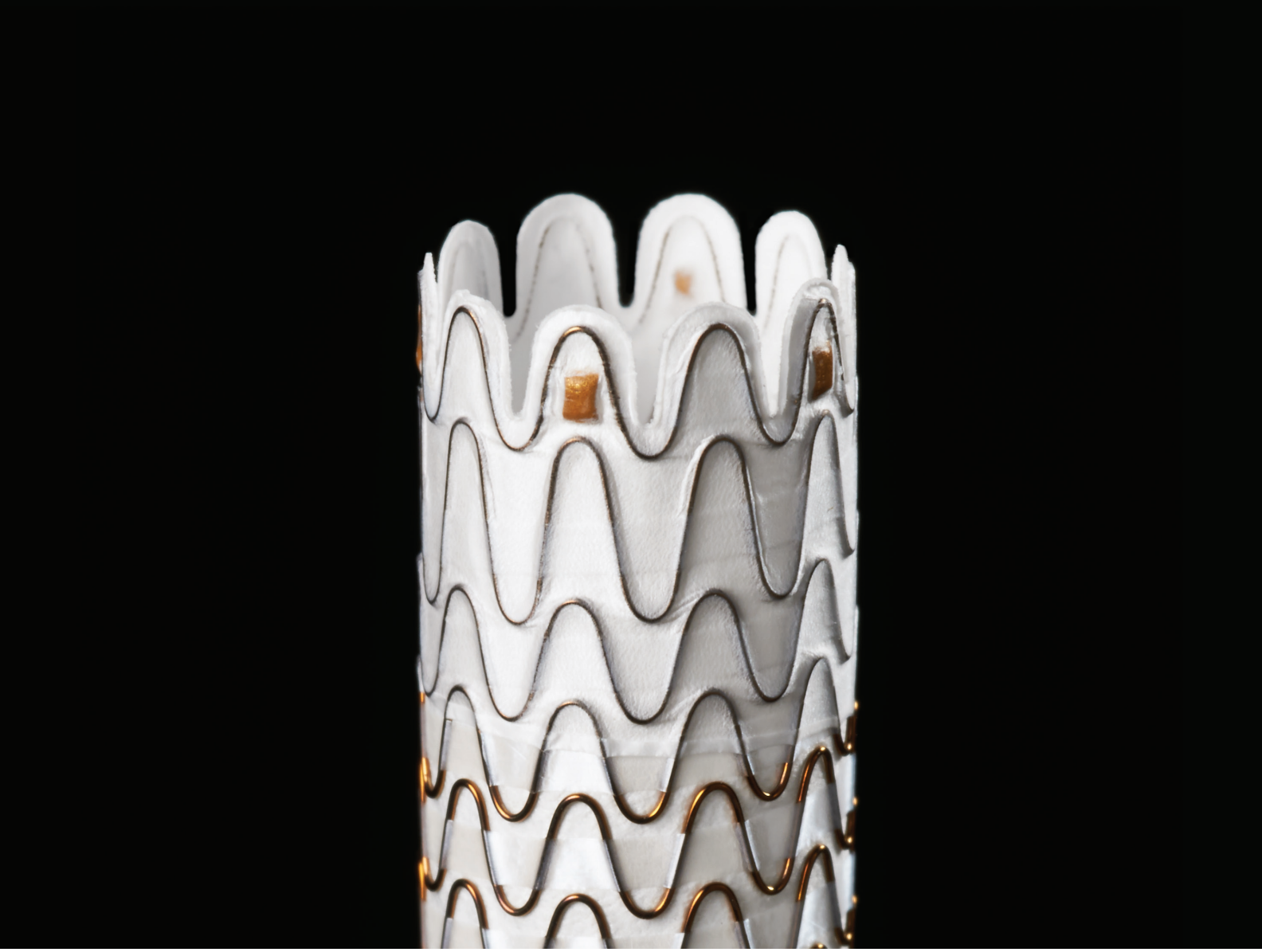

GORE® VIABAHN® Endoprosthesis with Heparin Bioactive Surface*

CONFIGURATIONS

* Heparin Bioactive Surface is synonymous with the CBAS Heparin Surface.

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Sizing table

0.035" Guidewire compatibility (with radiopaque markers)

Device sizing		Introducer sheath (Fr)						
Endoprosthesis labeled diameter* (mm)	Recommended vessel diameter† (mm)	2.5 cm Device length*	5 cm Device length*	7.5 cm Device length*	10 cm Device length*	15 cm Device length*	25 cm Device length*	Recommended balloon diameter for device touch-up‡ (mm)
5	4.0–4.7	7	7	7	7	7	7	5
6	4.8–5.5	7	7	7	7	7	7	6
7	5.6–6.5	8	8	8	8	8	8	7
8	6.6–7.5	8	8	8	8	8	8	8
9	7.6–8.5	–	8	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

0.014" or 0.018" Guidewire compatibility (with radiopaque markers)

Device sizing		Introducer sheath (Fr)						
Endoprosthesis labeled diameter* (mm)	Recommended vessel diameter† (mm)	2.5 cm Device length*	5 cm Device length*	7.5 cm Device length*	10 cm Device length*	15 cm Device length*	25 cm Device length*	Recommended balloon diameter for device touch-up‡ (mm)
5	4.0–4.7	6	6	6	6	6	6	5
6	4.8–5.5	6	6	6	6	6	6	6
7	5.6–6.5	7	7	7	7	7	7	7
8	6.6–7.5	7	7	7	7	7	7¶	8

Catalogue listings

0.035" Guidewire

Catalogue number	Endoprosthesis labeled diameter* (mm)	Endoprosthesis length* (cm)	Catheter length‖ (cm)	Recommended vessel diameter† (mm)	Device profile (Fr)
VBHR050202A	5	2.5	120	4.0–4.7	7
VBHR050502A	5	5	120	4.0–4.7	7
VBHR050702A	5	7.5	120	4.0–4.7	7
VBHR051002A	5	10	120	4.0–4.7	7
VBHR051502A	5	15	120	4.0–4.7	7
VBHR052502A	5	25	120	4.0–4.7	7
VBHR060201A	6	2.5	75	4.8–5.5	7
VBHR060202A	6	2.5	120	4.8–5.5	7
VBHR060501A	6	5	75	4.8–5.5	7
VBHR060502A	6	5	120	4.8–5.5	7
VBHR060702A	6	7.5	120	4.8–5.5	7
VBHR061001A	6	10	75	4.8–5.5	7

Catalogue number	Endoprosthesis labeled diameter* (mm)	Endoprosthesis length* (cm)	Catheter length‖ (cm)	Recommended vessel diameter† (mm)	Device profile (Fr)
VBHR061002A	6	10	120	4.8–5.5	7
VBHR061501A	6	15	75	4.8–5.5	7
VBHR061502A	6	15	120	4.8–5.5	7
VBHR062502A	6	25	120	4.8–5.5	7
VBHR070201A	7	2.5	75	5.6–6.5	8
VBHR070202A	7	2.5	120	5.6–6.5	8
VBHR070501A	7	5	75	5.6–6.5	8
VBHR070502A	7	5	120	5.6–6.5	8
VBHR070702A	7	7.5	120	5.6–6.5	8
VBHR071001A	7	10	75	5.6–6.5	8
VBHR071002A	7	10	120	5.6–6.5	8
VBHR071501A	7	15	75	5.6–6.5	8
VBHR071502A	7	15	120	5.6–6.5	8
VBHR072502A	7	25	120	5.6–6.5	8
VBHR080201A	8	2.5	75	6.6–7.5	8
VBHR080202A	8	2.5	120	6.6–7.5	8
VBHR080501A	8	5	75	6.6–7.5	8
VBHR080502A	8	5	120	6.6–7.5	8
VBHR080702A	8	7.5	120	6.6–7.5	8
VBHR081001A	8	10	75	6.6–7.5	8
VBHR081002A	8	10	120	6.6–7.5	8
VBHR081501A	8	15	75	6.6–7.5	8
VBHR081502A	8	15	120	6.6–7.5	8
VBHR082502A	8	25	120	6.6–7.5	8
VBHR090502A	9	5	120	7.6–8.5	8
VBHR090702A	9	7.5	120	7.6–8.5	8
VBHR091002A	9	10	120	7.6–8.5	8
VBHR091502A	9	15	120	7.6–8.5	8
VBHR100502A	10	5	120	8.6–9.5	8
VBHR101002A	10	10	120	8.6–9.5	8
VBHR101502A	10	15	120	8.6–9.5	8
VBHR110502A	11	5	120	9.6–10.5	10
VBHR111002A	11	10	120	9.6–10.5	10
VBHR130502A	13	5	120	10.6–12.0	10§
VBHR131002A	13	10	120	10.6–12.0	10§

Catalogue listings

0.014 / 0.018" Guidewire

Catalogue number	Endoprosthesis labeled diameter* (mm)	Endoprosthesis length* (cm)	Catheter length‖ (cm)	Recommended vessel diameter† (mm)	Device profile (Fr)
VBJR050202A	5	2.5	120	4.0–4.7	6
VBJR050502A	5	5	120	4.0–4.7	6
VBJR050702A	5	7.5	120	4.0–4.7	6
VBJR051002A	5	10	120	4.0–4.7	6
VBJR051502A	5	15	120	4.0–4.7	6
VBJR052502A	5	25	120	4.0–4.7	6
VBJR060202A	6	2.5	120	4.8–5.5	6
VBJR060502A	6	5	120	4.8–5.5	6
VBJR060702A	6	7.5	120	4.8–5.5	6
VBJR061002A	6	10	120	4.8–5.5	6
VBJR061502A	6	15	120	4.8–5.5	6
VBJR062502A	6	25	120	4.8–5.5	6
VBJR070202A	7	2.5	120	5.6–6.5	7
VBJR070502A	7	5	120	5.6–6.5	7
VBJR070702A	7	7.5	120	5.6–6.5	7
VBJR071002A	7	10	120	5.6–6.5	7
VBJR071502A	7	15	120	5.6–6.5	7
VBJR072502A	7	25	120	5.6–6.5	7
VBJR080202A	8	2.5	120	6.6–7.5	7
VBJR080502A	8	5	120	6.6–7.5	7
VBJR080702A	8	7.5	120	6.6–7.5	7
VBJR081002A	8	10	120	6.6–7.5	7
VBJR081502A	8	15	120	6.6–7.5	7
VBJR082502A	8	25	120	6.6–7.5	7¶

Consult Instructions for Use
eifu.goremedical.com

INTENDED USE / INDICATIONS: The GORE® VIABAHN® Endoprosthesis is indicated for improving blood flow in patients with symptomatic peripheral arterial disease in superficial femoral artery de novo and restenotic lesions up to 270 mm in length with reference vessel diameters ranging from 4.0 – 7.5 mm. The GORE® VIABAHN® Endoprosthesis is indicated for improving blood flow in patients with symptomatic peripheral arterial disease in superficial femoral artery in-stent restenotic lesions up to 270 mm in length with reference vessel diameters ranging from 4.0 – 6.5 mm. The GORE® VIABAHN® Endoprosthesis is indicated for improving blood flow in patients with symptomatic peripheral arterial disease in iliac artery lesions up to 80 mm in length with reference vessel diameters ranging from 4.0 – 12 mm. The GORE® VIABAHN® Endoprosthesis is also indicated for the treatment of stenosis or thrombotic occlusion at the venous anastomosis of synthetic arteriovenous (AV) access grafts.

CONTRAINDICATIONS: The GORE® VIABAHN® Endoprosthesis is contraindicated for non-compliant lesions where full expansion of an angioplasty balloon catheter was not achieved during pre-dilatation, or where lesions cannot be dilated sufficiently to allow passage of the delivery system. Do not use the GORE® VIABAHN® Endoprosthesis with Heparin Bioactive Surface 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

* 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.
¶ The 8 mm x 25 cm device is not compatible with the 7 Fr COOK® CHECK-FLO® Sheath.

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Products listed may not be available in all markets.

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