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all sizes
0.035" guidewire
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PERFORMANCE
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Indicated for Use in the Superficial Femoral and Iliac Arteries (5 – 8 mm diameters)

TIP to HUB Device Deployment – 0.035" Guidewire

CATALOGUE NUMBER	ENDOPROSTHESIS LABELED DIAMETER ¹ (mm)	ENDOPROSTHESIS LENGTH ¹ (cm)	CATHETER LENGTH ² (cm)	RECOMMENDED VESSEL DIAMETER ³ (mm)	DEVICE PROFILE (Fr)	RECOMMENDED BALLOON DIAMETER FOR DEVICE TOUCH-UP (mm)
VBC050202	5	2.5	120	4.0 – 4.7	7	5
VBC050502	5	5.0	120	4.0 – 4.7	7	5
VBC051002	5	10.0	120	4.0 – 4.7	7	5
VBC051502	5	15.0	120	4.0 – 4.7	7	5
VBC060202	6	2.5	120	4.8 – 5.5	7	6
VBC060501	6	5.0	75	4.8 – 5.5	7	6
VBC060502	6	5.0	120	4.8 – 5.5	7	6
VBC061001	6	10.0	75	4.8 – 5.5	7	6
VBC061002	6	10.0	120	4.8 – 5.5	7	6
VBC061501	6	15.0	75	4.8 – 5.5	7	6
VBC061502	6	15.0	120	4.8 – 5.5	7	6
VBC070202	7	2.5	120	5.6 – 6.5	8	7
VBC070501	7	5.0	75	5.6 – 6.5	8	7
VBC070502	7	5.0	120	5.6 – 6.5	8	7
VBC071001	7	10.0	75	5.6 – 6.5	8	7
VBC071002	7	10.0	120	5.6 – 6.5	8	7
VBC071501	7	15.0	75	5.6 – 6.5	8	7
VBC071502	7	15.0	120	5.6 – 6.5	8	7
VBC080202	8	2.5	120	6.6 – 7.5	8	8
VBC080501	8	5.0	75	6.6 – 7.5	8	8
VBC080502	8	5.0	120	6.6 – 7.5	8	8
VBC081001	8	10.0	75	6.6 – 7.5	8	8
VBC081002	8	10.0	120	6.6 – 7.5	8	8
VBC081501	8	15.0	75	6.6 – 7.5	8	8
VBC081502	8	15.0	120	6.6 – 7.5	8	8

¹ Labeled device diameters and lengths are nominal.

² Ensure the guidewire is the appropriate size (see *Instructions for Use*) and has a length at least twice that of the delivery catheter.

³ Recommended endoprosthesis compression within the vessel is approximately 5 – 20%.

Indicated for Use in the Iliac Artery (9 – 13 mm diameters)

TIP to HUB Device Deployment – 0.035" Guidewire

CATALOGUE NUMBER	ENDOPROSTHESIS LABELED DIAMETER ¹ (mm)	ENDOPROSTHESIS LENGTH ¹ (cm)	CATHETER LENGTH ² (cm)	RECOMMENDED VESSEL DIAMETER ³ (mm)	DEVICE PROFILE (Fr)	RECOMMENDED BALLOON DIAMETER FOR DEVICE TOUCH-UP ⁵ (mm)
VBC090502	9	5.0	120	7.6 – 8.5	9	9
VBC091002	9	10.0	120	7.6 – 8.5	9	9
VBC091502	9	15.0	120	7.6 – 8.5	9	9
VBC100202	10	2.5	120	8.6 – 9.5	11 ⁴	10
VBC100502	10	5.0	120	8.6 – 9.5	11 ⁴	10
VBC101002	10	10.0	120	8.6 – 9.5	11 ⁴	10
VBC101502	10	15.0	120	8.6 – 9.5	11 ⁴	10
VBC110202	11	2.5	120	9.6 – 10.5	11	12
VBC110502	11	5.0	120	9.6 – 10.5	11	12
VBC111002	11	10.0	120	9.6 – 10.5	11	12
VBC130202	13	2.5	120	10.6 – 12.0	12	14
VBC130502	13	5.0	120	10.6 – 12.0	12	14
VBC131002	13	10.0	120	10.6 – 12.0	12	14

¹ Labeled device diameters and lengths are nominal.

² Ensure the guidewire is the appropriate size (see *Instructions for Use*) and has a length at least twice that of the delivery catheter.

³ Recommended endoprosthesis compression within the vessel is approximately 5 – 20%.

⁴ 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.

⁵ For the 11 and 13 mm diameter devices, balloon inflation pressure should not exceed 8 atm.

 Consult Instructions for Use

INDICATIONS FOR USE: The GORE VIABAHN® Endoprosthesis is indicated for improving blood flow in patients with symptomatic peripheral arterial disease in superficial femoral artery lesions 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 iliac artery lesions with reference vessel diameters ranging from 4.0 – 12 mm. **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 incidence of Heparin-Induced Thrombocytopenia (HIT) type II. Refer to *Instructions for Use* at goremedical.com for a complete description of all warnings, precautions and adverse events. Rx Only



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