GORE® VIABAHN®
Endoprosthesis with
PROPATEN Bioactive Surface

Together, improving life

GORE® VIABAHN® ENDOPROSTHESIS WITH PROPATEN BIOACTIVE SURFACE* — CONFIGURATIONS

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

Sizing Table

0.035” Guidewire compatible

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

0.018” Guidewire compatible

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

0.018” Guidewire compatible

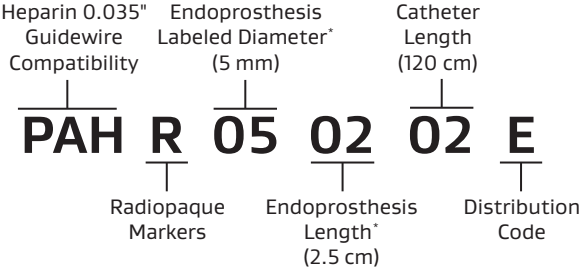
Catalogue Number	Endoprosthesis Labeled Diameter* (mm)	Endoprosthesis Length* (cm)	Catheter Length¶ (cm)	Recommended Vessel Diameter† (mm)	Device Profile (Fr)
PAJR050202E	5	2.5	120	4.0–4.7	6
PAJR050502E	5	5	120	4.0–4.7	6
PAJR050702E	5	7.5	120	4.0–4.7	6
PAJR051002E	5	10	120	4.0–4.7	6
PAJR051502E	5	15	120	4.0–4.7	6
PAJR052502E	5	25	120	4.0–4.7	6
PAJR060202E	6	2.5	120	4.8–5.5	6
PAJR060502E	6	5	120	4.8–5.5	6
PAJR060702E	6	7.5	120	4.8–5.5	6
PAJR061002E	6	10	120	4.8–5.5	6
PAJR061502E	6	15	120	4.8–5.5	6
PAJR062502E	6	25	120	4.8–5.5	6
PAJR070202E	7	2.5	120	5.6–6.5	7
PAJR070502E	7	5	120	5.6–6.5	7
PAJR070702E	7	7.5	120	5.6–6.5	7
PAJR071002E	7	10	120	5.6–6.5	7
PAJR071502E	7	15	120	5.6–6.5	7
PAJR072502E	7	25	120	5.6–6.5	7
PAJR080202E	8	2.5	120	6.6–7.5	7
PAJR080502E	8	5	120	6.6–7.5	7
PAJR080702E	8	7.5	120	6.6–7.5	7
PAJR081002E	8	10	120	6.6–7.5	7
PAJR081502E	8	15	120	6.6–7.5	7
PAJR082502E	8	25	120	6.6–7.5	7‖

Sizing Table

0.035” Guidewire compatible

Catalogue Number	Endoprosthesis Labeled Diameter* (mm)	Endoprosthesis Length* (cm)	Catheter Length¶ (cm)	Recommended Vessel Diameter† (mm)	Device Profile (Fr)
PAHR050202E	5	2.5	120	4.0–4.7	7
PAHR050502E	5	5	120	4.0–4.7	7
PAHR050702E	5	7.5	120	4.0–4.7	7
PAHR051002E	5	10	120	4.0–4.7	7
PAHR051502E	5	15	120	4.0–4.7	7
PAHR052502E	5	25	120	4.0–4.7	7
PAHR060201E	6	2.5	75	4.8–5.5	7
PAHR060202E	6	2.5	120	4.8–5.5	7
PAHR060501E	6	5	75	4.8–5.5	7
PAHR060502E	6	5	120	4.8–5.5	7
PAHR060702E	6	7.5	120	4.8–5.5	7
PAHR061001E	6	10	75	4.8–5.5	7
PAHR061002E	6	10	120	4.8–5.5	7
PAHR061501E	6	15	75	4.8–5.5	7
PAHR061502E	6	15	120	4.8–5.5	7
PAHR062502E	6	25	120	4.8–5.5	7
PAHR070201E	7	2.5	75	5.6–6.5	8
PAHR070202E	7	2.5	120	5.6–6.5	8
PAHR070501E	7	5	75	5.6–6.5	8
PAHR070502E	7	5	120	5.6–6.5	8
PAHR070702E	7	7.5	120	5.6–6.5	8
PAHR071001E	7	10	75	5.6–6.5	8
PAHR071002E	7	10	120	5.6–6.5	8
PAHR071501E	7	15	75	5.6–6.5	8
PAHR071502E	7	15	120	5.6–6.5	8
PAHR072502E	7	25	120	5.6–6.5	8
PAHR080201E	8	2.5	75	6.6–7.5	8
PAHR080202E	8	2.5	120	6.6–7.5	8
PAHR080501E	8	5	75	6.6–7.5	8
PAHR080502E	8	5	120	6.6–7.5	8
PAHR080702E	8	7.5	120	6.6–7.5	8
PAHR081001E	8	10	75	6.6–7.5	8
PAHR081002E	8	10	120	6.6–7.5	8
PAHR081501E	8	15	75	6.6–7.5	8
PAHR081502E	8	15	120	6.6–7.5	8
PAHR082502E	8	25	120	6.6–7.5	8
PAHR090502E	9	5	120	7.6–8.5	8
PAHR090702E	9	7.5	120	7.6–8.5	8
PAHR091002E	9	10	120	7.6–8.5	8
PAHR091502E	9	15	120	7.6–8.5	8
PAHR100502E	10	5	120	8.6–9.5	8
PAHR101002E	10	10	120	8.6–9.5	8
PAHR101502E	10	15	120	8.6–9.5	8
PAHR110502E	11	5	120	9.6–10.5	10
PAHR111002E	11	10	120	9.6–10.5	10
PAHR130502E	13	5	120	10.6–12.0	10§
PAHR131002E	13	10	120	10.6–12.0	10§

KEY: PRODUCT NUMBERING SYSTEM *(EXAMPLE ONLY)*



PAH Heparin 0.035" Guidewire Compatibility
PAJ Heparin 0.018" Guidewire Compatibility

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

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