

GORE® VIATORR® TIPS Endoprosthesis

DEVICE SPECIFICATIONS


GORE® VIATORR® TIPS Endoprosthesis

Designed for the Treatment of Portal Hypertension in Both De Novo and Revision Procedures

Catalogue number	Endoprosthesis internal diameter (mm)	Graft-lined length (cm)	Unlined length (cm)	Maximum guidewire diameter (in)	Hemostatic introducer sheath (Fr)	Maximum dilatation balloon diameter (mm)
PTB084275W	8	4	2	≤ 0.038	10	8
PTB085275W	8	5	2	≤ 0.038	10	8
PTB086275W	8	6	2	≤ 0.038	10	8
PTB087275W	8	7	2	≤ 0.038	10	8
PTB088275W	8	8	2	≤ 0.038	10	8
PTB104275W	10	4	2	≤ 0.038	10	10
PTB105275W	10	5	2	≤ 0.038	10	10
PTB106275W	10	6	2	≤ 0.038	10	10
PTB107275W	10	7	2	≤ 0.038	10	10
PTB108275W	10	8	2	≤ 0.038	10	10

Indications for Use: The GORE® VIATORR® TIPS Endoprosthesis is indicated for use in the de novo and revision treatment of portal hypertension and its complications such as variceal bleeding, gastropathy, refractory ascites, and / or hepatic hydrothorax.

 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.

GORE, VIATORR and designs are trademarks of W. L. Gore & Associates.
© 2020 W. L. Gore & Associates, Inc. AZ0872-EN1 MARCH 2020



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