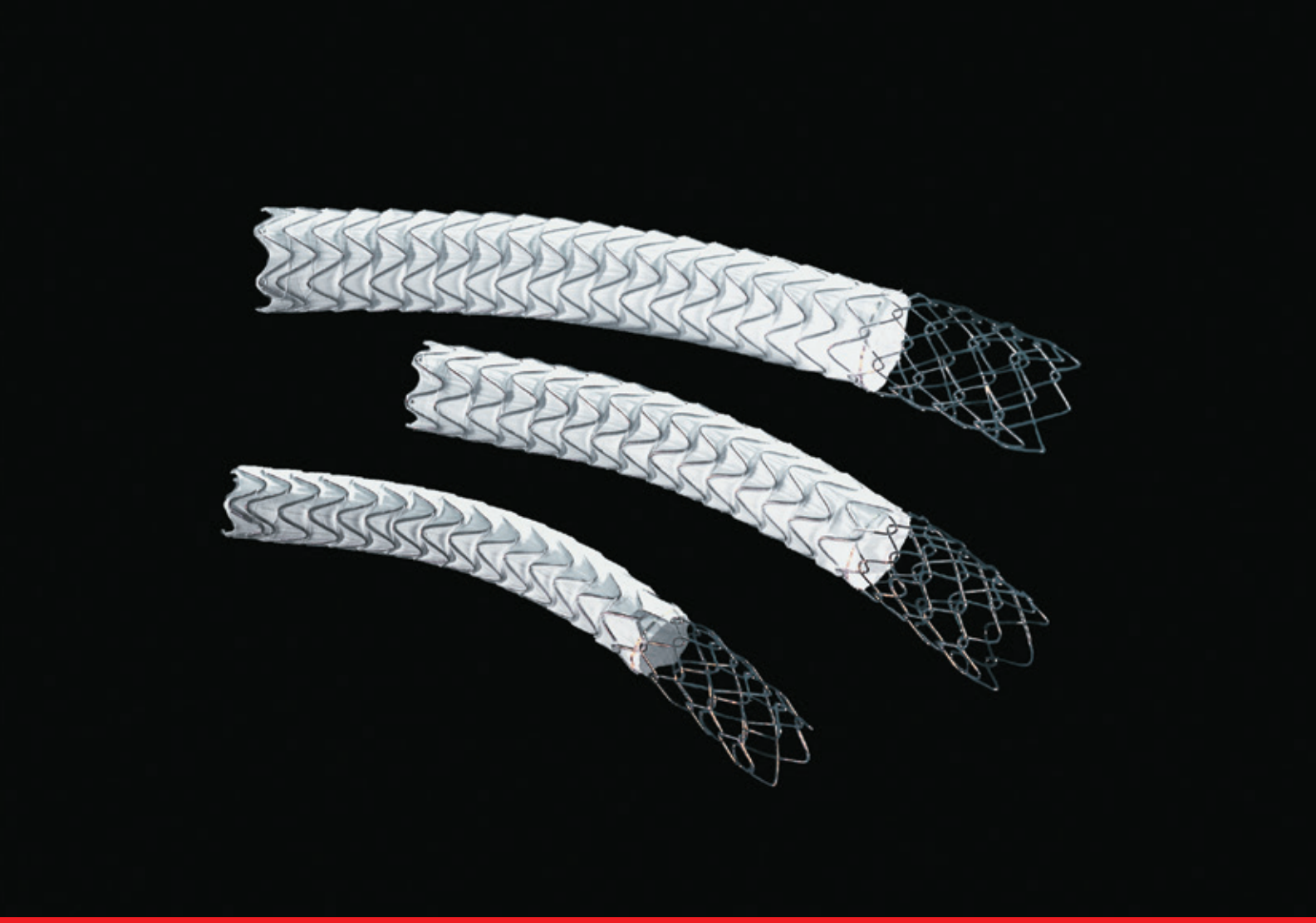

GORE® VIATORR® TIPS Endoprosthesis

DEVICE SPECIFICATIONS

GORE
Creative Technologies
Worldwide

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
Creative Technologies Worldwide

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