Reduce the risk of reintervention.

ASSURANCE YOU CAN COUNT ON.

ConMed

Authorized Distributor of Gore Medical Products

GORE VIABIL® BILIARY ENDOPROSTHESIS

Intended for palliation of malignant strictures in the biliary tree

UNIQUE ANTI-MIGRATION TECHNOLOGY aids in reducing the risk of reinterventions. The fully covered atraumatic anchoring fins securely hold the device within the duct, and

WE STAND BEHIND IT.

ANTI-MIGRATION ASSURANCE PROGRAM

Migrations are a known risk for any biliary endoprosthesis. With a 0.2% average reported migration rate, GORE® VIABIL® Biliary Endoprosthesis is the only fully covered metal stent with anti-migration technology proven to reduce the risk of reintervention. Gore is so confident in these outcomes, we are now offering a Device Replacement program for GORE® VIABIL® Biliary Endoprosthesis if a migration occurs.

PROGRAM DETAILS

Gore will provide a replacement device of identical dimensions for use with the patient whose device migrates within one year post implantation. The replacement device is only available if GORE® VIABIL® Biliary Endoprosthesis is implanted in accordance with the device *Instructions for Use* (The GORE® VIABIL® Biliary Endoprosthesis is intended for palliation of malignant strictures in the biliary tree) and the other terms of the program are satisfied. Replacement devices provided under this program are not eligible for the program. Claims under the program are limited to the replacement device. Upon receipt of the appropriate documentation, a replacement device will be provided pursuant to the program accompanied by a no-charge invoice shipped directly to the hospital. The hospital is responsible for reporting the no-charge replacement stent as a discount on the hospital's cost report. All reports of migration will be documented appropriately within the Gore internal product surveillance process and additional information may be requested. Migrations are a known risk of any biliary endoprosthesis. The provision of a replacement device as part of the program does not constitute an admission that there was a device malfunction or defect or that Gore, its employees or agents, or the Gore device caused or contributed to any complications or injuries. Please see the device *Instructions for Use* for further information on the device contraindications, warnings, precautions, and potential adverse events. The program is subject to modification or termination by W. L. Gore & Associates without prior notification and this program is only applicable for the United States.

Contact your ConMed Territory Manager for details of the program.

ConMed distributes the GORE® VIABIL® Biliary Endoprosthesis

You can also visit **goremedical.com/viabil** for more information.

GORE
Creative Technologies Worldwide

W. L. Gore & Associates, Inc.
Flagstaff, AZ 86004

+65.67332882 (Asia Pacific)
00800.6334.4673 (Europe)
800.437.8181 (United States)
928.779.2771 (United States)

goremedical.com

Data on file.

Refer to *Instructions for Use* for a complete description of all warnings, precautions, and contraindications. ℞ Only
Products listed may not be available in all markets.

MCM2017139