MRI Safety Information:

A patient with the GORE® VIABAHN® FORTEGRA Venous Stent may be safely scanned immediately after placement under the following conditions. Failure to follow these conditions may result in injury.
If information about a specific parameter is not included, there are no conditions associated with that parameter.

Device Name	GORE® VIABAHN® FORTEGRA Venous Stent
Static Magnetic Field Strength (B_0)	1.5T or 3.0T
Maximum Spatial Field Gradient	40 T/m (4,000 gauss/cm)
RF Excitation	There are no RF excitation restrictions
RF Transmit Coil Type	There are no Transmit Coil restrictions
Operating Mode	Normal Operating Mode
Maximum Whole-Body SAR	2 W/kg (Normal Operating Mode)
Scan Duration	2 W/kg whole-body averaged SAR for 60 minutes of continuous RF (a sequence or back to back series/scan without breaks)
MR Image Artifact	The presence of this implant may produce an image artifact. With a gradient echo pulse sequence in a 3.0T MR System, the artifact may extend up to 5mm from the implant. Under these conditions, the central portion of the lumen was visible.

For further information, visit
goremedical.com
800 528 8763

Consult Instructions for Use
eifu.goremedical.com

Caution

MR Conditional

30266000

FOR US ONLY
Patient Information Card

MD Medical Device

GORE® VIABAHN® FORTEGRA
Venous Stent

Patient Name

Date of Implant

Emergency Contact

Physician's Name

Physician's Phone

I am taking blood thinning medication(s): ☐ Yes ☐ No

Place product identification label in the space provided below.

GORE® VIABAHN® FORTEGRA
Venous Stent
REF
SN

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OCTOBER 2025
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