



LOWER PROFILE VBX STENT GRAFT

Same endoprosthesis and trusted performance,
now with greater versatility

**A lower delivery profile enables
a 1 Fr reduction on most sizes.**

Stent labeled/ nominal diameter (mm)	Stent length (mm)	Legacy profile (Fr)	Current profile (Fr)
5	15/19/29/39/59/79	7	6
6	15/19/29/39/59/79	7	6
7	15/19/29/39/59/79	7	6
8	29/39/59	7	7
8	79	8	7
8L	29/39	7	7
8L	59/79	8	8
9	29/39/59/79	8	7
10	29/39/59/79	8	7
11	29/39/59/79	8	8

6Fr
compatible
**GREATER
VERSATILITY**

All VBX Stent Graft configurations available in 0.035" over-the-wire 80 and 135 cm catheter lengths.

The **longest** balloon expandable stent graft is now 6 Fr compatible and still offers the **widest** range of diameter adjustability.¹⁻⁴

Stent labeled/nominal diameter (mm)	Maximum post-dilated stent diameter (mm) ^a
5	8
6	8
7	11
8	11
8L	16 ^b
9	13
10	13
11	16 ^b

All configurations available in 0.035" over-the-wire 80 and 135 cm catheter lengths.

a. Secondary balloon required to post-dilate the stent beyond its nominal deployed diameter (secondary balloon not included).

b. Technical limit of the device as determined by in vitro testing for the indicated use; device expansion beyond 13 mm was not studied as part of the Gore VBX FLEX Clinical Study. See *Instructions for Use*.

UP TO
79 mm
stent length¹



References

1. W. L. Gore & Associates. *GORE® VIABAHN® VBX Balloon Expandable Endoprosthesis Instructions for Use*. W. L. Gore & Associates; 2025. MD169334, Rev 8. Accessed April 28, 2026. <https://eifu.goremedical.com/>
2. BD. *The Control You Need to Deliver Accurate Treatment: LIFESTREAM™ Balloon Expandable Covered Stent*. BD; 2022. BD-52176v2. Accessed June 9, 2026. <https://www.bd.com/content/dam/bd-assets/na/peripheral-intervention/web-assets/us/documents/bd-52176-lifestream-br.pdf>
3. Atrium Medical Corporation. *iCAST Covered Stent System Instructions for Use*. Atrium Medical Corporation; 2025. EL012022, Rev AA.
4. U.S. Food and Drug Administration. *Summary of Safety and Effectiveness Data (SSED): LifeStream™ Balloon Expandable Vascular Covered Stent*. PMA P160024. Accessed June 2, 2026. https://www.accessdata.fda.gov/cdrh_docs/pdf16/P160024B.pdf

 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

INDICATIONS FOR USE IN THE US: The GORE® VIABAHN® VBX Balloon Expandable Endoprosthesis is indicated for the treatment of de novo or restenotic lesions found in iliac arteries with reference vessel diameters ranging from 5 mm–13 mm and lesion lengths up to 110 mm, including lesions at the aortic bifurcation. The GORE® VIABAHN® VBX Balloon Expandable Endoprosthesis is also indicated for use with thoracoabdominal and pararenal branched devices indicated with the GORE® VIABAHN® VBX Balloon Expandable Endoprosthesis as a branch component.

CONTRAINDICATIONS: Do not use the GORE® VIABAHN® VBX Balloon Expandable Endoprosthesis in patients with known hypersensitivity to heparin, including those patients who have had a previous incident of Heparin-Induced Thrombocytopenia (HIT) type II.

Products listed may not be available in all markets.

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W. L. Gore & Associates, Inc.
goremedical.com

Asia Pacific +65 67332882 Australia/New Zealand 1800 680 424 Europe 00800 6334 4673
United States Flagstaff, AZ 86004 800 437 8181 928 779 2771

