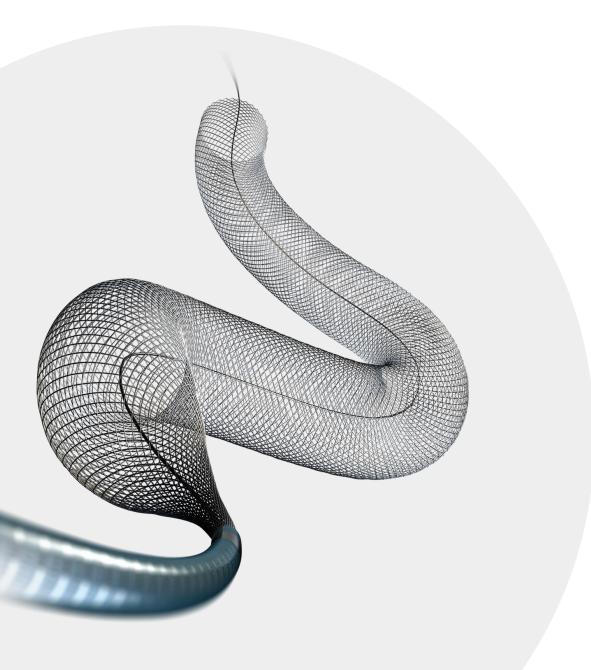
Pipeline[™] Vantage

Embolization device with shield technology™

The Vantage response

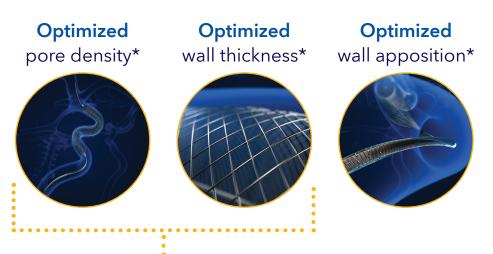


Reduced material thrombogenicty

By covalently bonding phosphorylcholine to the surface of the implant, Shield Technology[™] achieves a scientifically proven reduction in implant material thrombogenicity, thereby reducing blood contact activation in-vitro.¹⁻⁵

Promotes flow diversion effect

Endothelizalization is important to the success of flow diversion therapy. Pipeline[™] Vantage focuses on 3 key areas that promote the flow diversion effect.



Maintain proven metal to vessel coverage of $\mathsf{Pipeline}^{\scriptscriptstyle\mathsf{TM}}$

More control, precise delivery

Improvements to both the device and delivery system provides **more control** during navigation, deployment and resheathing so you can achieve a **precise delivery**, when treating aneurysms near and far.

Key design features*:

- Advanced Re-sheathing mechanism for **reduced friction**
- Single core wire for **increased pushability** and **stability**
- 44% thinner ePTFE sleeves for **faster** distal braid release
- Compatible with Phenom[™] 21
 Catheter for easier distal access
- Compatible with Rist[™] Radial Access
 Guide Catheter make the turn

4th Generation Pipeline™ Vantage with Shield Technology™ launched in Canada 4th Generation Pipeline[™] Vantage with Shield Technology™ launched in Europe Shield 1yr study <u>4</u>5) Premier study Pflex study <u>B</u> 6 PUFS 5 year follow-up results **CNS** award winner 2nd Generation Pipeline[™] Flex launched in Canada 2nd and 3rd Generation Pipeline[™] Flex & Shield Technology™ launched in Europe Intrepid study <u>\$</u> Results of PUFS 5 year study <u>\$</u>3 Hong Kong experience Ankara experience <u>\$</u> 1st Generation Pipeline[™] Device launched in Canada Australian registry <u>8</u> Pita study <u>}</u> **Buenos Aires experience** 1st Generation Pipeline[™] device launched in Europe

Start of PUFS 5 year study

<u>B</u>

Product catalog number	Diameter (mm)	Length (mm)
PED3-021-250-10	2.5	10
PED3-021-250-12	2.5	12
PED3-021-250-14	2.5	14
PED3-021-250-16	2.5	16
PED3-021-250-20	2.5	20
PED3-021-275-12	2.75	12
PED3-021-275-14	2.75	14
PED3-021-275-16	2.75	16
PED3-021-275-20	2.75	20
PED3-021-300-12	3.0	12
PED3-021-300-14	3.0	14
PED3-021-300-16	3.0	16
PED3-021-300-20	3.0	20
PED3-021-325-12	3.25	12
PED3-021-325-14	3.25	14
PED3-021-325-16	3.25	16
PED3-021-325-20	3.25	20
PED3-021-350-12	3.5	12
PED3-021-350-14	3.5	14
PED3-021-350-16	3.5	16
PED3-021-350-20	3.5	20
PED3-021-350-25	3.5	25
PED3-027-350-12	3.5	12
PED3-027-350-14	3.5	14
PED3-027-350-16	3.5	16
PED3-027-350-20	3.5	20
PED3-027-350-25	3.5	25

1. Medtronic Internal Study, D00422708 Rev. A , Competitive Test Report - Material Thrombogenicity Evaluation of Flow Diversion Devices.

2. Girdhar G, Li J, Kostousov L, Wainwright J, Chandler WL. In-vitro thrombogenicity assessment of flow diversion and aneurysm bridging devices. J Thromb Thrombolysis. 2015 Nov;40(4):437-43. doi: 10.1007/s11239-015-1228-0. PMID: 25975924.

3. D00422708 Rev A.

4. Girdhar G, Ubl S, Jahanbekam R, Thinamany S, Belu A, Wainwright J, Wolf MF. Thrombogenicity assessment of Pipeline, Pipeline Shield, Derivo and P64 flow diverters in an in vitro pulsatile flow human blood loop model. eNeurologicalSci. 2019 Jan 8;14:77-84.

5. Girdhar G, Andersen A, Pangerl E, Jahanbekam R, Ubl S, Nguyen K, Wainwright J, Wolf MF. Thrombogenicity assessment of Pipeline Flex, Pipeline Shield, and FRED flow diverters in an in vitro human blood physiological flow loop model. J Biomed Mater Res A. 2018 Dec; 106(12):3195-3202.

See the device manual for detailed information regarding the instructions for use, indications, contraindications, warnings, precautions, and potential adverse events. For further information, contact your local Medtronic representative and/or consult the Medtronic website at medtronic.ca The Pipeline™ Vantage Embolization Device with Shield Technology™ is intended for use with or without embolic coils for the treatment of wide neck intracranial aneurysms (defined as having a large neck greater than 4 mm and/or a dome-to-neck ratio less than or equal to 1.5) that are not amenable to treatment with surgical clipping. Patients should be 21 years of age or older.

Medtronic

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medtronic.ca/PipelineVantage

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