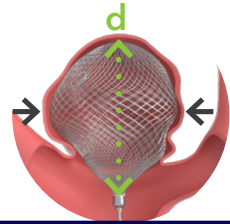
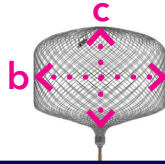
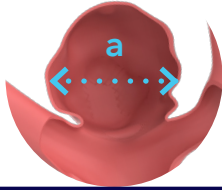


Artisite™ Intrasaccular Device

Sizing guide



Maximum average aneurysm width (mm) [a]	Artisite™ device width and height (mm) [b x c]	Compressed Device height when sized at max. avg. aneurysm width (mm) [d]
3.4	4.5 x 3.0	4.5
3.4	4.5 x 4.0	5.2
3.8	5.0 x 3.0	4.9
3.8	5.0 x 4.0	5.5
4.1	5.5 x 3.0	5.2
4.1	5.5 x 4.0	5.9
4.1	5.5 x 5.0	6.9
4.5	6.0 x 3.0	5.6
4.5	6.0 x 4.0	6.2
4.5	6.0 x 5.0	7.2
4.9	6.5 x 3.0	5.9
4.9	6.5 x 4.0	6.6
4.9	6.5 x 5.0	7.6
5.3	7.0 x 3.0	6.3
5.3	7.0 x 4.0	7.0
5.3	7.0 x 5.0	7.9
5.6	7.5 x 4.0	7.4
5.6	7.5 x 5.0	8.3
6.0	8.0 x 4.0	7.8
6.0	8.0 x 5.0	8.7

Note: If the measured average aneurysm width is between widths [a], target a larger group of devices while accounting for aneurysm height. Aneurysm height should be less than the compressed device height [d] for proper neck coverage.

This material should not be considered the exclusive source of information, it does not replace or supersede information contained in the device manual(s). Please note that the intended use of a product may vary depending on geographical approvals. See the device manual(s) for detailed information regarding the intended use, the (implant) procedure, indications, contraindications, warnings, precautions, and potential adverse events. For an MRI compatible device(s), consult the MRI information in the device manual(s) before performing a MRI. If a device is eligible for eIFU usage, instructions for use can be found at Medtronic's website manuals.medtronic.com Manuals can be viewed using a current version of any major internet browser. For best results, use Adobe Acrobat® Reader with the browser. Medtronic products placed on European markets comply with EU and UK legislation (if applicable) on medical devices. For any further information, contact your local Medtronic representative.

⌈
Estimated width oversizing factor
⌋

+1

+1.5

+2