stryker Neurovascular

Purposeful innovation in mind.

Bending expectations of conformability and stability.

Selection Guide

Product Number	Stent Diameter	Stent Length	Unconstrained Stent Diameter	Recommended Parent Vessel Diameter
M003EZAS30150	3.0mm	15mm	3.5mm	≥2.0 and <3.0mm
M003EZAS30210*	3.0mm	21mm	3.5mm	≥2.0 and <3.0mm
M003EZAS30240	3.0mm	24mm	3.5mm	≥2.0 and <3.0mm
M003EZAS40150	4.0mm	15mm	4.5mm	≥3.0 and <4.0mm
M003EZAS40240	4.0mm	24mm	4.5mm	≥3.0 and <4.0mm
M003EZAS40300*	4.0mm	30mm	4.5mm	≥3.0 and <4.0mm
M003EZAS45210	4.5mm	21mm	5.0mm	≥4.0 and ≤4.5mm
M003EZAS45300	4.5mm	30mm	5.0mm	≥4.0 and ≤4.5mm

All stent sizes are compatible with Stryker Neurovascular 0.0165-0.017in (0.42-0.43mm) ID microcatheters, excluding Tracker™ 17 Microcatheter.
The stents above are available without a distal delivery wire tip.
*These stent sizes will be available starting January, 2016.

- ***Adaptive cell structure***
  is designed to enhance stent opening and wall apposition
- ***Segmental™ Expansion***
  is designed to provide radial force and stability
- ***Low foreshortening***
  and predictable recrossability simplify use
- ***All stent sizes deliverable***
  through Excelsior™ SL-10™ Microcatheter

Neuroform Atlas™

STENT SYSTEM

See package insert for complete indications, contraindications, warnings and instructions for use.

INTENDED USE/INDICATIONS FOR USE
The Neuroform Atlas™ Stent System is intended to be used with occlusive devices in the treatment of intracranial aneurysms.

THIS DOCUMENT IS INTENDED SOLELY FOR THE USE OF HEALTHCARE PROFESSIONALS.
A physician must always rely on his or her own professional clinical judgment when deciding whether to use a particular product when treating a particular patient. Stryker does not dispense medical advice and recommends that physicians be trained in the use of any particular product before using it in a procedure. The information presented is intended to demonstrate the breadth of Stryker product offerings. A physician must always refer to the package insert, product label and/or instructions for use before using any Stryker product. Products may not be available in all markets because product availability is subject to the regulatory and/or medical practices in individual markets. Please contact your Stryker representative if you have questions about the availability of Stryker products in your area. The Stryker products listed above are CE marked according to the Medical Device Directive 93/42/EEC.

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