

NV PSR INSPIRE-A Pipeline™ Vantage Post Approval Study

NCT06604884

Status	RECRUITING
Sponsor	Medtronic Neurovascular Clinical Affairs
Enrollment	118 participants

Key Eligibility Criteria

Inclusion (3)

- Patient or legally authorized representative (LAR) has provided written informed consent using the Ethics Board and Medtronic approved Informed Consent Form and agrees to comply with the protocol requirements. HIPAA/data protection authorization has been provided and signed by the patient (or patient's LAR) as applicable per local law.
- Patient has an intracranial aneurysm intended to be treated with the Pipeline™ Vantage Embolization Device with Shield Technology™.
- Patient is an adult per local law at time of consent.

Exclusion (5)

- Patient with any contraindications for the device or procedure per the Pipeline™ Vantage Device local geography IFU.
- Patient who may be unable to complete the study follow-up.
- The Investigator determined that the health of the patient may be compromised by the patient's enrollment.
- Female patient who is known to be pregnant or is breastfeeding or wishes to become pregnant during participation in the study.
- Patient is currently enrolled in, or plans to enroll in, any concurrent drug/device study that may confound the study results.

Locations (4 total)

Wellstar Research Institute, Marietta, Georgia, United States
University of Iowa Hospitals and Clinics, Iowa City, Iowa, United States
McLaren Healthcare, Flint, Michigan, United States
... and 1 more locations