

CHOICE

Principal Investigator:

Safwan Kassas, MD (Covenant)

Sub-Investigators:

Rodney Diehl, DO

Ronald Bays, MD

Sponsor:

Abbott Vascular

Study Purpose:

Carotid Stenting For High Surgical-Risk Patients; Evaluating Outcomes Through The Collection Of Clinical Evidence

Objective:

The primary objective is to provide additional information that the commercially available Abbott Vascular Carotid Stent Systems and Embolic Protection Systems can be used successfully by a wide range of physicians under commercial use conditions. To provide an ongoing post-market surveillance mechanism for documentation of clinical outcomes and for possible extension of the Centers for Medicare and Medicaid Services (CMS) coverage to a broader group of patients.

Inclusion Criteria:

- ♥ Patient or patient's legally authorized representative provided informed consent.
- ♥ Patient is considered at high risk for carotid endarterectomy (CEA).
- ♥ Patient requires percutaneous carotid angioplasty and stenting for carotid artery disease.
- ♥ Patients physician intends to use an RX Acculink with the RX Accunet in the carotid artery or an Xact with the Emboshield or Emboshield Nav6 in the carotid artery as per the FDA approved Indications for Use as outlined.

Exclusion Criteria:

There are no exclusion criteria for this study

Status:

Active enrollment

If you have any questions, please feel free to contact the coordinators, and they will be happy to answer any questions you have regarding this study.