

EDUCATE

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Sponsor:

Medtronic Vascular

Study Purpose:

The purpose of this study is to collect additional safety information on the Endeavor stent, which has already been approved by the Food and Drug Administration (FDA).

Objective:

Study subjects with ischemic heart disease due to stenotic lesions in either native coronary arteries or coronary artery bypass grafts (CABG) undergoing PCI with stent placement are eligible to be enrolled in this study. Subjects may have multi-vessel treatment. Subjects eligible for randomization into the CPI should not have contraindications to prolonged DAPT noted at the time of randomization.

Inclusion Criteria:

1. Patient is > 18 years of age.
2. The patient or patient's legal representative has consented to participate and has authorized the collection and release of his medical information by signing the "Subject Informed Consent Form".
3. Patients undergoing percutaneous intervention with Endeavor stent deployment (or has within 24 hours).
4. The patient is willing and able to cooperate with study procedures and required follow up visits.

Exclusion Criteria:

1. Index procedure stent placement with stent diameter < 2.5 mm or > 3.5 mm.

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2. Pregnant women.
3. Current medical condition with a life expectancy of less than 3 years.
4. The patient is currently participating in another investigational device or drug study that clinically interferes with EDUCATE. The patient may only be enrolled in EDUCATE once.
5. Patients with hypersensitivity or allergies to one of the drugs or components indicated in the Instructions for Use.
6. Patients in whom antiplatelet and/or anticoagulation therapy is contraindicated.
7. Patients treated with any stent other than the Endeavor stent during the index procedure.

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.