Serrantor OCT Study

NCT06434194

Status	RECRUITING
Phase	Not Applicable
Sponsor	Cagent Vascular LLC
Enrollment	60 participants

Key Eligibility Criteria

Inclusion (3)

- Rutherford clinical category 4-6 of the target limb
- Age of subjects is \>18 years old
- Patients has given informed consent to participate in this study

Exclusion (4)

- De novo or restenotic (without prior stent) stenosis (e70%) or occlusion
- Target lesion is in the BTK arteries, including below the knee popliteal, tibioperoneal trunk, tibial, peroneal, and pedal arteries.
- Angiographic visual estimated reference vessel diameter is between 2.0 and 5.0 mm.
- Lesion length less than 220 mm

Locations (2 total)

Columbia University, New York, New York, United States
Weill Cornell Medical, New York, New York, United States