

Evaluation of the Clinical Outcomes of the Embolic Protection System in Preventing Distal Embolism During Femoropopliteal Debulking Procedures

NCT07282301

Status RECRUITING
Sponsor RenJi Hospital
Enrollment 150 participants

Key Eligibility Criteria

Inclusion (9)

- Age \geq 18 years
- Rutherford classification 2-5
- Patients with chronic femoropopliteal obstruction or thromboembolism who require debulking prior to other endovascular treatments.
- Patients who understand the purpose of this study, participate in the study voluntarily, sign the informed consent and are willing to receive follow-up visits
- The guidewire needs to pass through the lesion
- ... and 4 more (see full listing online)

Exclusion (7)

- Patients with stroke, cerebral hemorrhage, gastrointestinal hemorrhage, or cardiac infarction within 3 months prior to enrollment
- Patients with known allergy to heparin, aspirin, other antiplatelet drugs, contrast agents, etc.
- Patients who have been treated with drugs that interfere with this clinical trial or intraoperatively with other special vascular bed preparation devices, such as plaque volume reduction devices, special balloons, etc., within the last 3 months;
- Pregnant and lactating women
- Patients who are unable or unwilling to participate in this trial
- ... and 2 more (see full listing online)

Locations (1 total)

Renji Hospital, Shanghai, Shanghai Municipality, China