TriClip Japan Post-Approval Study

NCT07509658

Status RECRUITING
Sponsor Abbott Medical Devices
Enrollment 250 participants

Key Eligibility Criteria

Inclusion (2)

- Subjects are eligible to receive the TriClip System per the current approved indications for use (IFU)
- Subjects provide written informed consent prior to conducting any investigation-specific procedures not considered standard of care.

Exclusion (1)

- Please note: It is not recommended that subjects enrolled in the study participate in any other therapeutic clinical study.

Locations (1 total)

National Cerebral & Cardiovascular Center Hospital, Suita, Osaka, Japan