Post-market Clinical Follow-up Data Collection From Procedures With BIOTRONIK EP Products

NCT05560958

Status	RECRUITING
Sponsor	Biotronik SE & Co. KG
Enrollment	280 participants

Key Eligibility Criteria

Inclusion (9)

- Indication for diagnostic or therapeutic EP intervention
- EP intervention is planned to involve the use of BIOTRONIK EP products from at least 2 of the 3 following categories:
- BIOTRONIK catheter (AlCath, ViaCath, MultiCath, Khelix)
- BIOTRONIK external device (Qubic Stim, Qubic RF, Qiona)
- BIOTRONIK transseptal sheath (Senovo Bi-Flex)

... and 4 more (see full listing online)

Exclusion (4)

- Age less than 18 years
- Pregnant or breastfeeding
- Prior participation in this study with performed EP procedure
- Participation in an interventional clinical investigation in parallel to the BIO\|COLLECT.EP study

Locations (1 total)

Städtisches Krankenhaus Friedrichshafen, Friedrichshafen, Germany