

Efficacy and Safety of Micra AV2 Transcatheter Pacing System

NCT07356505

Status	RECRUITING
Phase	Not Applicable
Sponsor	Universitaire Ziekenhuizen KU Leuven
Enrollment	150 participants

Key Eligibility Criteria

Inclusion (5)

- Voluntary written informed consent of the participant or their legally authorized representative has been obtained prior to any screening procedures
- At least 18 years of age at the time of signing the Informed Consent Form (ICF)
- Participant will be implanted with a Micra AV2 for an approved indication for use
- Participant with a history of any type of high-degree AV block
- Participant is willing and able to comply the study protocol

Exclusion (1)

- Subject implanted with a Micra AV2 on a non-permanent basis (e.g. CIED infection) 2. Subject is pregnant (if required by local law, women of child-bearing potential must undergo a pregnancy test within seven days prior to Micra AV2 implant procedure) 3. Subject currently enrolled or planning to participate in a potentially confounding drug or device trial during the study.

Locations (1 total)

University Hospitals of Leuven, Leuven, Flemish Brabant, Belgium