Aveir VR Coverage With Evidence Development Post-Approval Study

NCT05336877

Status RECRUITING
Sponsor Abbott Medical Devices
Enrollment 8,744 participants

Key Eligibility Criteria

Inclusion (4)

- The study cohort will include all Medicare patients with continuous claims data implanted with an Aveir VR leadless pacemaker or a full-system single-chamber ventricular transvenous pacemaker (from any manufacturer) in any US location
- Medicare beneficiaries implanted with an Aveir VR leadless pacemaker on or after the study start date (i.e., the date of Aveir VR market approval) will be included in the study.
- OR
- Medicare beneficiaries implanted with a full system (e.g. lead and generator) single-chamber ventricular transvenous pacemaker on or after the study start date

Exclusion (1)

- None

Locations (1 total)

Abbott, Sylmar, California, United States

DISCLAIMER: This document is for informational purposes only and does not constitute medical advice. Always consult your healthcare provider before enrolling in any clinical trial. Information may not be up to date — verify details at ClinicalTrials.gov. Generated by ClinicalTrialsFinder.org.