Lifetech AcuMark™ Sizing Balloon Post-Market Clinical Follow-up Study

NCT06700174

Status	RECRUITING
Sponsor	Lifetech Scientific (Shenzhen) Co., Ltd.
Enrollment	100 participants

Key Eligibility Criteria

Inclusion (4)

- Patients who have been confirmed with single secundum atrial septal defect (ASD) by echocardiography;
- The anatomy of ASD is suitable for percutaneous closure;
- Patients are scheduled to perform ASD closure;
- Patients or legally authorized representative(s) who are willing and capable of providing informed consent.

Exclusion (4)

- Any contraindication for ASD closure;
- Patients have ostium primum ASD or coronary sinus ASD;
- Patients who are pregnant or breastfeeding;
- Patient is currently participating in another clinical trial that has not yet completed its primary endpoint.

Locations (5 total)

Ankara (Bilkent) City Hospital, Ankara, Turkey (Türkiye)
Gazi Yasargil Woman and Child Hospital, Diyarbak1r, Turkey (Türkiye)
Gaziantep University Hospital, Gaziantep, Turkey (Türkiye)
... and 2 more locations