Follow-up Study to Monitor the Efficacy and Safety of the Occlutech® mVSD Cases

NCT05329350

Status	RECRUITING
Sponsor	Occlutech International AB
Enrollment	50 participants

Key Eligibility Criteria

Inclusion (13)

- A participant will be eligible for study participation if he/she meets the indication and area of application as laid down in the IFU. The Occlutech mVSD is indicated for percutaneous occlusion of hemodynamically or clinically significant muscular ventricular septal defects.
- Any age
- Male or female.
- Patients understand the nature of the study and provide their informed consent to participation.
- Patients willing and able to attend the follow-up visits and procedures foreseen by study CIP.

... and 8 more (see full listing online)

Exclusion (15)

- The device is contraindicated for participants known to have any of the following:
- Active bacterial infections
- Active infection at the time of implantation
- Allergy to antiplatelet or anticoagulant therapy
- Allergy to nickel and/or titanium and/or nickel/titanium-based materials

... and 10 more (see full listing online)

Locations (5 total)

Children's Health Ireland, Dublin, Ireland
0zmir ^ehir Hastanesi, Izmir, Bayrakl1, Turkey (Türkiye)
Eski_ehir Osmangazi Üniversite Hastanesi, Eski_ehir, Odunpazar1, Turkey (Türkiye)
... and 2 more locations