

Evaluation of the Effectiveness of Two Methods of Regional Anesthesia During S-ICD Implantation Procedure.

NCT06520150

Status	RECRUITING
Phase	Not Applicable
Sponsor	4th Military Clinical Hospital with Polyclinic, Poland
Enrollment	32 participants

Key Eligibility Criteria

Inclusion (2)

- patients aged 18-90 years requiring S-ICD implantation for cardiac indications,
- high risk of anesthesia (ASA II or higher), patients giving informed consent to participate in the study.

Exclusion (4)

- absolute contraindications to regional anesthesia such as generalized coagulation disorders, inflammation at the site of anesthesia, allergy to the drugs used for regional anesthesia
- lack of consent of the patient for the conducted procedure,
- inability to cooperate with the patient - such as severe mental retardation, unregulated mental illness with motor agitation, advanced neurodegenerative diseases such as Parkinson's disease/Alzheimer's,
- inability to give informed consent to participate in the study.

Locations (1 total)

4th Military Clinical Hospital with Polyclinic, Wroclaw, Lower Silesian Voivodeship, Poland