

Transvaginal Human Acellular Dermal Matrix for Prolapse Treatment

NCT06404931

Status	RECRUITING
Phase	Phase 2
Sponsor	Hospital Mutua de Terrassa
Enrollment	90 participants

Key Eligibility Criteria

Inclusion (5)

- Women with symptomatic POP with indication of vaginal surgery.
- Women with hiatal ballooning or levator ani injury evaluated by 3D pelvic floor ultrasound.
- Women e18 years old.
- Women wishing to complete a 36-month follow-up.
- Understand and accept the study procedures and sign the informed consent.

Exclusion (5)

- Women with previous pelvic floor or prolapse surgery
- Women with POP grade IV
- Women with chronic pelvic pain.
- Not being able to understand the nature of the study and/or the procedures to be followed.
- Not signing the informed consent

Locations (1 total)

Hospital Mutua de Terrassa, Terrassa, Spain