Effectiveness of Radiofrequency Combined With Ultrasound for the Treatment of Postpartum Dyspareunia

NCT07504991

Status	RECRUITING
Phase	Not Applicable
Sponsor	Institut Investigacio Sanitaria Pere Virgili
Enrollment	34 participants

Key Eligibility Criteria

Inclusion (1)

- \- The study will enroll women of childbearing age with a medical history of at least one prior vaginal delivery. Participants must report persistent pain during penetrative sexual intercourse, meeting the clinical criteria for dyspareunia. Pain intensity will be assessed using the validated Visual Analogue Scale (VAS) ranging from 0 to 10. Eligible patients must demonstrate a moderate to severe pain level, defined as a VAS score of 4 or higher. Inclusion is independent of the time elapsed between the last delivery and the current clinical diagnosis

Exclusion (2)

- Contraindications to the study treatment: Women with any medical condition that precludes the use of the study technology
- Non-postpartum dyspareunia

Locations (1 total)

Centro Procrear, Tarragona, TARRAGONA, Spain