Sacral Neuromodulation for Chronic Pelvic Pain

NCT06150599

Status	RECRUITING
Phase	Not Applicable
Sponsor	Corewell Health East
Enrollment	25 participants

Key Eligibility Criteria

Inclusion (8)

- Provision of signed and dated informed consent form
- Stated willingness to comply with all study procedures and availability for the duration of the study
- Female, aged 22-70
- Chronic pelvic pain (= pain below umbilicus) score of 4 or greater on 10-point VAS, present for 6 months or greater (screening patient)
- Failed at least 1 or more conservative treatments (e.g. pelvic floor physical therapy, biofeedback, behavioral modification, oral pharmacotherapy, bladder instillations)

... and 3 more (see full listing online)

Exclusion (16)

- History of any active pelvic cancer
- Any significant medical condition that is likely to interfere with study procedures, device operation, or likely to confound evaluation of study endpoints
- Concurrent pain management strategies within the past 3 months that may interfere or mask study intervention (e.g. pelvic floor physical therapy, shockwave therapy, trigger point injections, bladder instillations)
- Any psychiatric or personality disorder at the discretion of the study physician. This does not include depression or generalized anxiety disorders that are stable.
- Current symptomatic urinary tract infection (UTI) or more than 6 UTIs in past year

... and 11 more (see full listing online)

Locations (1 total)

Corewell Health William Beaumont University Hospital, Royal Oak, Michigan, United States