EPPIC: Easing Pelvic Pain Interventions Clinical Research Program

NCT05127616

Status	RECRUITING
Phase	Not Applicable
Sponsor	State University of New York at Buffalo
Enrollment	240 participants

Key Eligibility Criteria

Inclusion (15)

- Ages 18-70 years (inclusive)
- Male or female
- All genders, races, ethnic groups
- MD-confirmed diagnosis of IC/BPS or CP/CPPS by study urologist or urogynecologist
- Pelvic pain including uncomfortable sensations of pressure or discomfort that are not described as outright pain) of at least six months duration

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

Exclusion (13)

- Presence of a neurological condition (e.g., MS, Parkinson's disease, paraplegia) affecting the bladder
- The presence of a symptomatic urethral stricture (males only)
- History of cystitis caused by tuberculosis or radiation or chemotherapies
- Participant has been diagnosed and treated for a pelvic-related malignancy (colon, bladder, prostate, ovarian, endometrial, uterine, testicular, penile, cervical, vaginal, or rectal cancer)
- Participant has a medical condition(s) whose nature or severity (unstable, life threatening, etc.) would influence adversely the conduct of the clinical trial, confound interpretation of study results, and/or compromise volunteer safety and engagement with study demands.

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

Locations (3 total)

UCLA, Los Angeles, California, United States
University of Michigan, Ann Arbor, Michigan, United States
University at Buffalo (the only clinical site where treatment is delivered), Buffalo, New York, United States