

Effectiveness of oral Solifenacin versus minimally invasive surgery in treatment of urge incontinence.

ACTRN12617000284369

Status	RECRUITING
Sponsor	Mugla Sitki Kocman University
Enrollment	192 participants

Plain Language Summary

This study is comparing medication (solifenacin, a bladder tablet) with a minimally invasive surgical procedure for women who experience urgent urinary leakage (urge incontinence) at least twice a day. The surgery tightens specific ligaments that support the bladder, while the medication works to calm an overactive bladder. Both treatments are being tested to see which works better.

You may be eligible if:

- You are a woman aged 21 or older
- You experience urge incontinence with leakage at least twice per day on an average day
- You have a mild to moderate bladder prolapse (POPQ grade 1–3)
- Your urge symptoms improve when a speculum is gently inserted into the vagina

You may NOT be eligible if:

- You have a 3rd or 4th degree uterine prolapse
- You have a urinary tract infection
- You have a known neurological condition such as multiple sclerosis
- You have previously had vaginal surgery (other than a sling procedure or hysterectomy)

Talk to your doctor about whether this trial might be right for you.

Key Eligibility Criteria

Inclusion (3)

- Urge incontinence with wetting twice or more per on an average day
- POPQ grade 1-3 cystocele or apical POP
- Improvement of urge symptoms by gentle insertion of speculum into the posterior vaginal fornix

Exclusion (4)

- Patients with 3rd or 4th degree uterine POP
- Urinary tract infections
- Known neurological conditions, for example multiple sclerosis
- Patients with previous vaginal surgery (other than midurethral sling or hysterectomy).

Locations (1 total)

Mugla, Turkey

<https://www.anzctr.org.au/Trial/Registration/TrialReview.aspx?ACTRN=ACTRN12617000284369>

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