Women with UI

Screening for eligibility (week -1)

Baseline assessment (week 0)
- 1-hour pad test
- 72-hour bladder diary
- ICIQ-UI Short Form

500 women with pure SUI

Central randomization

Treatment group (n=250)
EA, 3 sessions per week for 6 weeks
(Weeks 1-6)

Control group (n=250)
Sham EA, 3 sessions per week for 6 weeks
(Weeks 1-6)

24-week follow-up (Weeks 7-30)

Outcome measures

- Baseline
- Week 2
- Week 6
- Week 18
- Week 30

- Change in amount of urine leakage at week 6 from baseline measured by 1-hour pad test;
- 72-hour IEF (based on 72-hour bladder diary);
- ICIQ-UI Short Form score;
- Severity of urinary incontinence (based on 72-hour bladder diary);
- Self-report assessment of therapeutic effect (no baseline);
- Weekly consumption of pads;
- Application of other treatments for SUI;
- Subgroup analysis stratified by incontinence severity

Evaluation of safety, discomfort and acceptance of EA (at weeks 2 and 6)

Data collection and analysis