

**Non-surgical treatment for women with stress urinary incontinence (SUI)****Study eligibility form**

Assessor initials: \_\_\_\_\_

Date assessed: \_\_\_\_\_

**Study identifier**

(surname of first author + year of publication)

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|  |
|--|

**Type of study**

Q1. Is the study a randomised or quasi-randomised trial?  
(quasi-randomised = alternation, day of week, etc.)

|               |         |                |
|---------------|---------|----------------|
| Yes           | Unclear | No             |
| ↓             | ↓       | ↓              |
| Go to         |         |                |
| next question |         | <b>Exclude</b> |

**Participants in the study**

Q2. Are some or all of the participants in the study adult women with stress urinary incontinence, mixed urinary incontinence (with stress as predominant pattern), or undiagnosed or not-characterised urinary incontinence?

|               |         |                |
|---------------|---------|----------------|
| Yes           | Unclear | No             |
| ↓             | ↓       | ↓              |
| Go to         |         |                |
| next question |         | <b>Exclude</b> |

NB. For studies recruiting men and women, data must be reported separately for women.

**Interventions in the study**

Q3. Does the study involve at least one of the following interventions?

Lifestyle, pelvic floor muscle training ± biofeedback, vaginal cones, electrical stimulation (not nerve stimulation), electromagnetic stimulation, vaginal cones, behavioural therapy (e.g. bladder training), serotonin and nonrepinephrine reuptake inhibitors (SNRI), injectables, mechanical devices, containment/absorbent pads, catheters

|               |         |                |
|---------------|---------|----------------|
| Yes           | Unclear | No             |
| ↓             | ↓       | ↓              |
| Go to         |         |                |
| next question |         | <b>Exclude</b> |

**Outcomes in the study**

Q4. Does the study report one or more of the following outcomes?

Cure/improvement rates, quantification of symptoms, quality of life

|                |         |                |
|----------------|---------|----------------|
| Yes            | Unclear | No             |
| ↓              | ↓       | ↓              |
| <b>Include</b> |         | <b>Exclude</b> |

Final decision (subject to clarification of 'unclear' points)

**Include   Unclear   Exclude**

*Write here if the study is relevant for updating Cochrane reviews* ☞