

# Stage 3 Pilot Study: a decision tool to improve the management of urinary incontinence in women in the community

<b>Submission date</b> 18/03/2011	<b>Recruitment status</b> No longer recruiting	<input checked="" type="checkbox"/> Prospectively registered
<b>Registration date</b> 18/03/2011	<b>Overall study status</b> Completed	<input type="checkbox"/> Protocol
<b>Last Edited</b> 04/07/2017	<b>Condition category</b> Urological and Genital Diseases	<input type="checkbox"/> Statistical analysis plan
		<input type="checkbox"/> Results
		<input type="checkbox"/> Individual participant data
		<input type="checkbox"/> Record updated in last year

## Plain English summary of protocol

Not provided at time of registration

## Contact information

### Type(s)

Scientific

### Contact name

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## Additional identifiers

### Protocol serial number

9940

## Study information

### Scientific Title

Stage 3 Pilot Study: a decision tool to improve the management of urinary incontinence in women in the community

**Study objectives**

The overall aim of the project is to develop a decision tool for the management of urinary incontinence in women in the community that can be used by health care professionals based in primary and community care settings (GPs, practice and community nurses). The decision tool has been developed with input from clinicians through a series of focus groups and is being tested for validity and reliability. The aim of the current pilot study is the implementation of the decision tool in practice.

**Ethics approval required**

Old ethics approval format

**Ethics approval(s)**

11/H1307/1

**Study design**

Both; Interventional; Design type: Diagnosis, Process of Care, Treatment

**Primary study design**

Interventional

**Study type(s)**

Treatment

**Health condition(s) or problem(s) studied**

Topic: Primary Care Research Network for England, Renal and Urogenital; Subtopic: Not Assigned, Renal and Urogenital (all Subtopics); Disease: All Diseases, Urogenital

**Interventions**

We seek to recruit at least 60 health care professionals from each of the two study sites and as many patients as present with symptoms of UI during the course of the pilot study.

Use of the decision tool: The decision tool has been developed to improve the management of urinary incontinence in women in the community.

Follow Up Length: 4 month(s); Study Entry : Other; Details: Patients will not be randomised. It is the health care professionals who will be randomised.

**Intervention Type**

Other

**Phase**

Not Applicable

**Primary outcome(s)**

Amelioration of symptoms of urinary incontinence; Timepoint(s): Baseline, 1 months, 4 months

**Key secondary outcome(s)**

Not provided at the time of registration

**Completion date**

31/07/2011

## Eligibility

### Key inclusion criteria

1. Healthcare professionals: Any healthcare professional who works in a community/primary care setting whose role involves assessment and management of urinary incontinence in women in the community. Health care professionals will be working within two primary care trusts where NHS permissions will be sought for the study to proceed
2. Patients: Women who are over 18 years of age, not pregnant and have symptoms of urinary incontinence are eligible to be invited into the study.; Target Gender: Male & Female ; Lower Age Limit 18 no age limit or unit specified

### Participant type(s)

Patient

### Healthy volunteers allowed

No

### Age group

Adult

### Lower age limit

18 years

### Sex

All

### Key exclusion criteria

1. Health care professionals who were involved in the intial development of the decision tool
2. Patients who are under 18 years of age
3. Patients who are pregnant
4. Patients who are unable to give informed consent

### Date of first enrolment

01/04/2011

### Date of final enrolment

31/07/2011

## Locations

### Countries of recruitment

United Kingdom

England

### Study participating centre

**University of Leeds**  
Leeds  
United Kingdom  
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## Sponsor information

**Organisation**  
University of York (UK)

**ROR**  
<https://ror.org/04m01e293>

## Funder(s)

**Funder type**  
Government

**Funder Name**  
National Institute for Health Research (NIHR) (UK) - Research for Patient Benefit Programme  
(ref: grant codes: PB-PG-1207-15081)

## Results and Publications

**Individual participant data (IPD) sharing plan**

**IPD sharing plan summary**  
Not provided at time of registration

### Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
<a href="#">Participant information sheet</a>	Participant information sheet	11/11/2025	11/11/2025	No	Yes