

# Evaluation of services for urinary dysfunction

<b>Submission date</b> 25/10/2000	<b>Recruitment status</b> No longer recruiting	<input type="checkbox"/> Prospectively registered
<b>Registration date</b> 25/10/2000	<b>Overall study status</b> Completed	<input type="checkbox"/> Protocol
<b>Last Edited</b> 12/09/2007	<b>Condition category</b> Urological and Genital Diseases	<input type="checkbox"/> Statistical analysis plan
		<input checked="" type="checkbox"/> Results
		<input type="checkbox"/> Individual participant data

**Plain English summary of protocol**  
Not provided at time of registration

## Contact information

**Type(s)**  
Scientific

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## Additional identifiers

**Protocol serial number**  
G9410491

## Study information

**Scientific Title**

### Study objectives

The objective of the phase I trial is to compare a new Continence Nurse Practitioner (CNP) led service with existing service provision for individuals with incontinence and lower urinary tract symptoms in terms of symptom severity, associated factors, impact and economic implications.

Treatment in the nursing service lasts eight weeks. The new nursing service delivers a package of evidence based interventions for the provision of continence care which are protocol driven.

**Ethics approval required**

Old ethics approval format

**Ethics approval(s)**

Not provided at time of registration

**Study design**

Randomised controlled trial

**Primary study design**

Intentional

**Study type(s)**

Not Specified

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

Urinary dysfunction

**Interventions**

1. Existing service provision: general practitioners and primary health care team
2. Nurse led service provision: new service instituting evidence based assessment and interventions provided by specially trained nurses

**Intervention Type**

Other

**Phase**

Not Specified

**Primary outcome(s)**

An independent interview is carried out 13 weeks and 26 weeks after randomisation in both arms of the trial . This measures symptom severity, associated factors, impact (social and psychological) and economic aspects of the condition.

**Key secondary outcome(s)**

Not provided at time of registration

**Completion date**

31/03/2002

**Eligibility****Key inclusion criteria**

Urinary dysfunction measured on a postal questionnaire (incontinence, nocturia, frequent voiding, urinary urgency)

**Participant type(s)**

Patient

**Healthy volunteers allowed**

No

**Age group**

Adult

**Sex**

All

**Key exclusion criteria**

1. Pregnancy
2. Malignancy
3. Fistula
4. Those already in receipt of treatment

**Date of first enrolment**

01/06/1996

**Date of final enrolment**

31/03/2002

**Locations**

**Countries of recruitment**

United Kingdom

England

**Study participating centre**

Department of Epidemiology and Public Health

Leicester

United Kingdom

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**Sponsor information**

**Organisation**

Medical Research Council (MRC) (UK)

**Funder(s)**

**Funder type**

Research council

**Funder Name**

Medical Research Council (UK)

**Alternative Name(s)**

Medical Research Council (United Kingdom), UK Medical Research Council, Medical Research Committee and Advisory Council, MRC

**Funding Body Type**

Government organisation

**Funding Body Subtype**

National government

**Location**

United Kingdom

## 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="#">Results article</a>	Results	01/08/2002		Yes	No