

The measurement of urinary incontinence via survey questionnaires

Submission date 10/10/2005	Recruitment status No longer recruiting	☒ Prospectively registered
		☐ Protocol
Registration date 13/10/2005	Overall study status Completed	☐ Statistical analysis plan
		☐ Results
Last Edited 19/10/2009	Condition category Urological and Genital Diseases	☐ Individual participant data
		☐ Record updated in last year

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

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Additional identifiers

Study information

Scientific Title

Study objectives

The trial aims to see if lead-in questions will affect the reported rate of urinary incontinence

Ethics approval required

Old ethics approval format

Ethics approval(s)

Not provided at time of registration

Primary study design

Interventional

Study design

Randomised controlled trial

Study type(s)

Other

Health condition(s) or problem(s) studied

Urinary incontinence

Interventions

The control group will be asked:

Do you ever leak any water (urine) when you don't mean to? Yes/No

The intervention group will be asked some related questions preceding the urinary incontinence question i.e.

1. Do you have to pass water (urine) at least once during the night? Yes/No

2. Do you have to pass water at least five times per day? Yes/No

3. Do you need to pass water when you hear running water? Yes/No

4. Do you sometimes have the sensation of not emptying your bladder after you have finished passing water? Yes/No

5. Do you get a sudden strong urge to pass water? Yes/No

6. Do you have difficulty in controlling your water? Yes/No

7. Do you ever leak any water when you don't mean to? Yes/No

Differences in the response rate between the two groups will be analysed for statistical significance.

Intervention Type

Other

Phase

Not Specified

Primary outcome(s)

Information on the effect of leading questions on urinary incontinence.

Key secondary outcome(s)

The survey also contains questions relating to a range of health and lifestyle issues e.g smoking, alcohol use and diet.

Completion date

25/11/2005

Eligibility

Key inclusion criteria

1. Patients registered with Bury General Practitioners.
2. People over the age of 18.

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Adult

Lower age limit

18 Years

Sex

All

Key exclusion criteria

People under the age of 18.

Date of first enrolment

17/10/2005

Date of final enrolment

25/11/2005

Locations

Countries of recruitment

United Kingdom

England

Study participating centre

Public Health Department

Bury

United Kingdom

BL9 0EN

Sponsor information

Organisation

Bury Primary Care Trust (UK)

Funder(s)**Funder type**

Government

Funder Name

Funding provided by the public health department at Bury Primary Care Trust (UK)

Results and Publications**Individual participant data (IPD) sharing plan****IPD sharing plan summary**

Not provided at time of registration