The measurement of urinary incontinence via survey questionnaires

Submission date	Recruitment status	[X] Prospectively registered
10/10/2005	No longer recruiting	Protocol
Registration date	Overall study status	Statistical analysis plan
13/10/2005	Completed	Results
Last Edited	Condition category	Individual participant data
19/10/2009	Urological and Genital Diseases	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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Contact details

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

N/A

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

Study design

Randomised controlled trial

Primary study design

Interventional

Secondary study design

Randomised controlled trial

Study setting(s)

GP practice

Study type(s)

Other

Participant information sheet

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 measure

Information on the effect of leading questions on urinary incontinence.

Secondary outcome measures

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

Overall study start date

17/10/2005

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

Age group

Adult

Lower age limit

18 Years

Sex

Both

Target number of participants

1430

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

England

United Kingdom

Study participating centre Public Health Department

Bury United Kingdom BL9 0EN

Sponsor information

Organisation

Bury Primary Care Trust (UK)

Sponsor details

21 Silver Street
Bury
England
United Kingdom
BL9 0EN
+44 (0)161 762 3136
paul.campbell@burypct.nhs.uk

Sponsor type

Hospital/treatment centre

Website

http://www.burypct.nhs.uk

Funder(s)

Funder type

Government

Funder Name

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

Results and Publications

Publication and dissemination planNot provided at time of registration

Intention to publish date

Individual participant data (IPD) sharing plan

IPD sharing plan summaryNot provided at time of registration