Reduction of anxiety before urodynamics: a randomised trial of pre-test education

Submission date	Recruitment status	Prospectively registered
29/09/2006	No longer recruiting	☐ Protocol
Registration date	Overall study status	Statistical analysis plan
29/09/2006	Completed	☐ Results
Last Edited	Condition category	Individual participant data
20/02/2020	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

Dr A Doshani

Contact details

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers N0123173227

Study information

Scientific Title

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Study objectives

The study aims to compare between the effect of written information and a guided tour on anxiety levels in women attending the urodynamic clinics.

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)

Other

Study type(s)

Treatment

Participant information sheet

Not available in web format, please use the contact details below to request a patient information sheet

Health condition(s) or problem(s) studied

Urological and Genital Diseases

Interventions

Randomised controlled trial. Clinical Study, Human Physiology, Inpatient Admission, Multicentre Study: National, NHS Patients, Study of Non-Therapeutic Procedure.

Written information and a guided tour vs standard practice.

Intervention Type

Other

Phase

Not Specified

Primary outcome measure

The difference in STAI score between women in the intervention and control group.

Secondary outcome measures

Not provided at time of registration

Overall study start date

16/09/2005

Completion date

01/12/2005

Eligibility

Key inclusion criteria

150 questionnaires will be given to all women attending the urodynamics clinic.

Participant type(s)

Patient

Age group

Not Specified

Sex

Female

Target number of participants

150

Key exclusion criteria

Not provided at time of registration

Date of first enrolment

16/09/2005

Date of final enrolment

01/12/2005

Locations

Countries of recruitment

England

United Kingdom

Study participating centre University Hospitals of Leicester

Leicester United Kingdom LE1 4PW

Sponsor information

Organisation

Record Provided by the NHSTCT Register - 2006 Update - Department of Health

Sponsor details

The Department of Health, Richmond House, 79 Whitehall London United Kingdom SW1A 2NL +44 (0)20 7307 2622 dhmail@doh.gsi.org.uk

Sponsor type

Government

Website

http://www.dh.gov.uk/Home/fs/en

Funder(s)

Funder type

Government

Funder Name

University Hospitals of Leicester NHS Trust (UK), Department Research Funds

Results and Publications

Publication and dissemination plan

Not provided at time of registration

Intention to publish date

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?

Other publications scoping study 01/12/2009 Yes No