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A comparison between the use of transurethral resection of the prostate (TURP) with bipolar cutting loop diathermy for the treatment of benign prostatic hypertrophy

Submission date 30/09/2005	**Recruitment status** No longer recruiting	[_] Prospectively registered [_] Protocol [_] Statistical analysis plan [_] Results [_] Individual participant data [_] Record updated in last year
Registration date 30/09/2005	**Overall study status** Completed	
Last Edited 14/03/2014	**Condition category** Urological and Genital Diseases	

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers
N0234156276

Study information

Scientific Title

Study objectives
Effects of standard TURP using 5% glycine as irrigation with bipolar resection which can use normal saline as irrigation fluid.

Ethics approval required
Old ethics approval format

Ethics approval(s)
Not provided at time of registration

Study design
Prospective randomised controlled trial

Primary study design
Interventional

Secondary study design
Randomised controlled trial

Study setting(s)
Not specified

Study type(s)
Not Specified

Participant information sheet

Health condition(s) or problem(s) studied
Urological and Genital Diseases: Benign prostatic hyperplasia (BPH)

Interventions
Randomised to either bipolar resection with normal saline or TURP with glycine

Intervention Type
Other

Phase
Not Specified

Primary outcome measure
ECG evidence of ischaemia, chest pain, cardiac arrhythmias, neurological symptoms, biochemistry results, haematology research

Secondary outcome measures

Not provided at time of registration

Overall study start date
01/12/2004

Completion date
01/12/2006

Eligibility

Key inclusion criteria
All patients scheduled for TURP under spinal anaesthesia will be approached.

Participant type(s)
Patient

Age group
Not Specified

Sex
Male

Target number of participants
210

Key exclusion criteria
Inability to give informed consent, general anaesthesia

Date of first enrolment
01/12/2004

Date of final enrolment
01/12/2006

Locations

Countries of recruitment
England

United Kingdom

Study participating centre
BUI
Bristol
United Kingdom
BS16 1ND

Sponsor information

Organisation
Department of Health

Sponsor details
Richmond House
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Sponsor type
Government

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

Funder(s)

Funder type
Government

Funder Name
North Bristol NHS Trust (UK), NHS R&D Support Funding

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