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A prospective randomised double blind and placebo controlled trial of tamsulosin for post operative urinary retention in men undergoing hip and knee arthroplasty and spinal surgery

Submission date 12/09/2003	**Recruitment status** No longer recruiting	[_] Prospectively registered [_] Protocol
Registration date 12/09/2003	**Overall study status** Completed	[_] Statistical analysis plan [_] Results
Last Edited 09/03/2018	**Condition category** Signs and Symptoms	[_] 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

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

N0084096576

Study information

Scientific Title
A prospective randomised double blind and placebo controlled trial of tamsulosin for post operative urinary retention in men undergoing hip and knee arthroplasty and spinal surgery

Study objectives
Can perioperative use of tamsulosin reduce the incidence of postoperative urinary retention at statistically and clinically significant levels?

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)
Hospital

Study type(s)
Treatment

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

Health condition(s) or problem(s) studied
Post operative urinary retention

Interventions
Randomised controlled trial comparing (a) Tamsulosin and (b) placebo.

Intervention Type
Drug

Phase
Not Applicable

Drug/device/biological/vaccine name(s)

Tamulosin

Primary outcome measure
Not provided at time of registration

Secondary outcome measures
Not provided at time of registration

Overall study start date
30/12/1999

Completion date
14/07/2004

Eligibility

Key inclusion criteria
512 patients in total

Participant type(s)
Patient

Age group
Adult

Sex
Male

Target number of participants
512

Key exclusion criteria
Does not match inclusion criteria

Date of first enrolment
30/12/1999

Date of final enrolment
14/07/2004

Locations

Countries of recruitment
England

United Kingdom

Study participating centre

Hull Royal Infirmary
Hull
United Kingdom
HU3 2JZ

Sponsor information

Organisation
Department of Health (UK)

Sponsor details
Richmond House
79 Whitehall
London
United Kingdom
SW1A 2NL

Sponsor type
Government

Website
http://www.doh.gov.uk

Funder(s)

Funder type
Government

Funder Name
The North and South Bank Research and Development Consortium (NHS R&D Support Funding)

Funder Name
British Orthopaedic Society

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