

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	<input type="checkbox"/> Prospectively registered
		<input type="checkbox"/> Protocol
Registration date 12/09/2003	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan
		<input type="checkbox"/> Results
Last Edited 09/03/2018	Condition category Signs and Symptoms	<input type="checkbox"/> Individual participant data
		<input type="checkbox"/> 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

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