

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

Protocol serial number

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

Study type(s)

Treatment

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(s)

Not provided at time of registration

Key secondary outcome(s))

Not provided at time of registration

Completion date

14/07/2004

Eligibility**Key inclusion criteria**

512 patients in total

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Adult

Sex

Male

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**

United Kingdom

England

Study participating centre

Hull Royal Infirmary

Hull

United Kingdom

HU3 2JZ

Sponsor information**Organisation**

Department of Health (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

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?
Participant information sheet	Participant information sheet	11/11/2025	11/11/2025	No	Yes