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Prospective double-blind randomised controlled trial to evaluate the role of Tamsulosin in the prevention of post-operative urinary retention following inguinal hernia repair

Submission date 30/09/2004	**Recruitment status** No longer recruiting	[ ] Prospectively registered [ ] Protocol
Registration date 30/09/2004	**Overall study status** Completed	[ ] Statistical analysis plan [ ] Results
Last Edited 25/03/2020	**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
N0186133788

Study information

Scientific Title
Prospective double-blind randomised controlled trial to evaluate the role of Tamsulosin in the prevention of post-operative urinary retention following inguinal hernia repair

Study objectives
Evaluation of the role of the alpha adrenergic blocker Tamsulosin in the prevention of post-operative urinary retention following inguinal hernia repair

Ethics approval required
Old ethics approval format

Ethics approval(s)
Not provided at time of registration

Study design
Prospective double-blind randomised controlled trial

Primary study design
Interventional

Secondary study design
Randomised controlled trial

Study setting(s)
Hospital

Study type(s)
Treatment

Participant information sheet

Health condition(s) or problem(s) studied
Signs and Symptoms: Post-operative urinary retention

Interventions
Tamsulosin vs standard practice

Intervention Type
Drug

Phase
Not Specified

Drug/device/biological/vaccine name(s)
Tamsulosin

Primary outcome measure
Not provided at time of registration

Secondary outcome measures
Not provided at time of registration

Overall study start date
22/07/2003

Completion date
21/01/2004

Eligibility

Key inclusion criteria
Not provided at time of registration

Participant type(s)
Patient

Age group
Not Specified

Sex
Not Specified

Target number of participants
200 patients

Key exclusion criteria
Not provided at time of registration

Date of first enrolment
22/07/2003

Date of final enrolment
21/01/2004

Locations

Countries of recruitment
England

United Kingdom

Study participating centre

Poole Hospital NHS Trust
Poole
United Kingdom
BH15 2JD

Sponsor information

Organisation
Department of Health

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

Sponsor type
Government

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

Funder(s)

Funder type
Government

Funder Name
Poole Hospital NHS Trust (UK)

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