

Randomised, controlled trial of the stoppa repair versus the Lichtenstein method for the repair of bilateral inguinal hernia: short-term outcome

Submission date 12/09/2003	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered
Registration date 12/09/2003	Overall study status Completed	<input type="checkbox"/> Protocol
Last Edited 14/04/2015	Condition category Urological and Genital Diseases	<input type="checkbox"/> Statistical analysis plan
		<input type="checkbox"/> Results
		<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

M0010058197

Study information

Scientific Title

Randomised, controlled trial of the stoppa repair versus the Lichtenstein method for the repair of bilateral inguinal hernia: short-term outcome

Study objectives

Post-operative complications and return to activity following bilateral inguinal hernia repair.

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

Health condition(s) or problem(s) studied

Urological and Genital Diseases: Inguinal hernia

Interventions

Stoppa repair versus Lichtenstein repair

Intervention Type

Other

Phase

Not Specified

Primary outcome measure

1. Post-operative complications
2. Post-operative pain
3. Return to normal activity

Secondary outcome measures

Not provided at time of registration

Overall study start date

01/05/1999

Completion date

01/05/2006

Eligibility

Key inclusion criteria

50 patients in each group, undergoing repair of bilateral inguinal hernia

Participant type(s)

Patient

Age group

Not Specified

Sex

Not Specified

Target number of participants

100

Key exclusion criteria

Not provided at time of registration

Date of first enrolment

01/05/1999

Date of final enrolment

01/05/2006

Locations

Countries of recruitment

England

United Kingdom

Study participating centre

Royal Bolton Hospital
Bolton
United Kingdom
BL4 0JR

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

Bolton Hospitals 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