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Prospective randomised trial to evaluate laparoscopic versus open bilayer patch herniorrhaphy for the treatment of primary unilateral inguinal hernia

Submission date 30/09/2004	**Recruitment status** No longer recruiting	[_] Prospectively registered [_] Protocol [_] Statistical analysis plan [_] Results [_] Individual participant data [_] Record updated in last year
Registration date 30/09/2004	**Overall study status** Completed	
Last Edited 01/05/2015	**Condition category** Urological and Genital Diseases	

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
N0084125082

Study information

Scientific Title
Prospective randomised trial to evaluate laparoscopic versus open bilayer patch herniorrhaphy for the treatment of primary unilateral inguinal hernia

Study objectives
To evaluate laparoscopic herniorrhaphy under general anaesthetic compared to open patch herniorrhaphy under local anaesthetic in the treatment of primary inguinal hernia.

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
Laparoscopic herniorrhaphy under general anaesthetic compared to open patch herniorrhaphy under local anaesthetic in the treatment of primary inguinal hernia.

Intervention Type
Other

Phase
Not Specified

Primary outcome measure
Not provided at time of registration

Secondary outcome measures
Not provided at time of registration

Overall study start date
29/05/2003

Completion date
01/06/2005

Eligibility

Key inclusion criteria
Not provided at time of registration

Participant type(s)
Patient

Age group
Adult

Sex
Both

Target number of participants
398 patients will be enrolled

Key exclusion criteria
Not provided at time of registration

Date of first enrolment
29/05/2003

Date of final enrolment
01/06/2005

Locations

Countries of recruitment
England

United Kingdom

Study participating centre
Hull Royal Infirmary
Hull
United Kingdom
HU3 2JZ

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
The North and South Bank Research and Development Consortium (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