Prospective randomised double blind trial of ilioinguinal nerve block versus local anaesthetic wound instillation in patients undergoing open inguinal herniorrhaphy

Submission date	Recruitment status	Prospectively registered
30/09/2005	No longer recruiting	☐ Protocol
Registration date	Overall study status	Statistical analysis plan
30/09/2005	Completed	Results
Last Edited	Condition category	Individual participant data
25/04/2014	Surgery	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

N0170118241

Study information

Scientific Title

Study objectives

Is infiltration of the operative wound with local anaesthetic as effective as an inguinal nerve block?

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)

Not specified

Study type(s)

Not Specified

Participant information sheet

Health condition(s) or problem(s) studied

Surgery: Open inquinal herniorrhaphy

Interventions

We will audit the use of local anaesthetic in an ilioinguinal nerve block as a local anaesthetic for pain at the end of inguinal hernia, and the alternative technique of instilling local anaesthetic into the operative field at the end of surgery.

Intervention Type

Procedure/Surgery

Phase

Not Specified

Primary outcome measure

To obtain objective information of pain relief after hernia surgery by instilling local anaesthetic and to compare the results with ilioinquinal nerve blockade.

Secondary outcome measures

Not provided at time of registration

Overall study start date

11/09/2002

Completion date

31/12/2003

Eligibility

Key inclusion criteria

A total of 60 patients, undergoing open inguinal herniorrhaphy. Male patients listed for open repair as day case of a primary unilateral inguinal hernia under the care of two NCH consultants.

Participant type(s)

Patient

Age group

Not Specified

Sex

Male

Target number of participants

60

Key exclusion criteria

No known exclusion criteria.

Date of first enrolment

11/09/2002

Date of final enrolment

31/12/2003

Locations

Countries of recruitment

England

United Kingdom

Study participating centre

Nottingham City Hospital

Nottingham United Kingdom NG5 1PB

Sponsor information

Organisation

Department of Health

Sponsor details

Richmond House 79 Whitehall London United Kingdom SW1A 2NL +44 (0)20 7307 2622 dhmail@doh.gsi.org.uk

Sponsor type

Government

Website

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

Funder(s)

Funder type

Government

Funder Name

Nottingham City Hospital NHS Trust (UK), NCH Directorate R&D Budget

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 summaryNot provided at time of registration