

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

Submission date 30/09/2005	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered
		<input type="checkbox"/> Protocol
Registration date 30/09/2005	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan
		<input type="checkbox"/> Results
Last Edited 25/04/2014	Condition category Surgery	<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

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

Primary study design

Interventional

Study design

Randomised controlled trial

Study type(s)

Not Specified

Health condition(s) or problem(s) studied

Surgery: Open inguinal 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(s)

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

Key secondary outcome(s)

Not provided at time of registration

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

Healthy volunteers allowed

No

Age group

Not Specified

Sex

Male

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**

United Kingdom

England

Study participating centre

Nottingham City Hospital

Nottingham

United Kingdom

NG5 1PB

Sponsor information**Organisation**

Department of Health

Funder(s)**Funder type**

Government

Funder Name

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

Results and Publications

Individual participant data (IPD) sharing plan

IPD sharing plan summary

Not provided at time of registration