

Study of effect of gabapentin on persistent pain after open inguinal hernia repair

Submission date 12/09/2003	Recruitment status Stopped	<input type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
Registration date 12/09/2003	Overall study status Stopped	<input type="checkbox"/> Statistical analysis plan <input type="checkbox"/> Results
Last Edited 18/10/2016	Condition category Digestive System	<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

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers
N0547083244

Study information

Scientific Title

Study of effect of gabapentin on persistent pain after open inguinal hernia repair

Study objectives

To establish whether gabapentin at a dose of 900-1800 mg per day is effective at reducing persistent pain after open 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

Not available in web format, please use contact details to request a participant information sheet

Health condition(s) or problem(s) studied

Inguinal hernia

Interventions

Randomised controlled trial: gabapentin at a dose of 900 - 1800 mg per day.

Intervention Type

Drug

Phase

Not Applicable

Drug/device/biological/vaccine name(s)

Gabapentin

Primary outcome measure

Not provided at time of registration

Secondary outcome measures

Not provided at time of registration

Overall study start date

31/05/2000

Completion date

31/05/2003

Eligibility

Key inclusion criteria

16 subjects, 16 controls

Participant type(s)

Patient

Age group

Adult

Sex

Both

Target number of participants

32

Key exclusion criteria

Not provided at time of registration

Date of first enrolment

31/05/2000

Date of final enrolment

31/05/2003

Locations

Countries of recruitment

England

United Kingdom

Study participating centre

West Norwich Hospital
Norwich
United Kingdom
NR2 3TU

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

Norfolk and Norwich University Hospital/Norwich Primary Care Trust (PCT) (UK) - East Norfolk and Waveney Research Consortium

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