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

Submission date 12/09/2003	**Recruitment status** Stopped	[ ] Prospectively registered [ ] Protocol [ ] Statistical analysis plan [ ] Results [ ] Individual participant data [ ] Record updated in last year
Registration date 12/09/2003	**Overall study status** Stopped	
Last Edited 18/10/2016	**Condition category** Digestive System	

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