

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

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

**Protocol serial number**  
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

**Study type(s)**

Treatment

**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(s)**

Not provided at time of registration

**Key secondary outcome(s)**

Not provided at time of registration

**Completion date**

31/05/2003

**Eligibility****Key inclusion criteria**

16 subjects, 16 controls

**Participant type(s)**

Patient

**Healthy volunteers allowed**

No

**Age group**

Adult

**Sex**

All

**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**

United Kingdom

England

**Study participating centre**

**West Norwich Hospital**

Norwich

United Kingdom

NR2 3TU

**Sponsor information****Organisation**

Department of Health (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**

**Individual participant data (IPD) sharing plan**

**IPD sharing plan summary**

Not provided at time of registration