

# Randomised double blind placebo controlled trial of the efficacy of gabapentin in improving analgesia in patients undergoing hysterectomy receiving patients controlled analgesia

<b>Submission date</b> 30/09/2005	<b>Recruitment status</b> No longer recruiting	<input type="checkbox"/> Prospectively registered
		<input type="checkbox"/> Protocol
<b>Registration date</b> 30/09/2005	<b>Overall study status</b> Completed	<input type="checkbox"/> Statistical analysis plan
		<input type="checkbox"/> Results
<b>Last Edited</b> 02/11/2016	<b>Condition category</b> Signs and Symptoms	<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

Dr B Brandner

### Contact details

Anaesthetics

Department of Anaesthesia

The Middlesex Hospital

Mortimer Street

London

United Kingdom

W1N 8AA

+44

brigitta.brandner@uclh.nhs.uk

## Additional identifiers

### Protocol serial number

N0263156073

# Study information

## Scientific Title

Randomised double blind placebo controlled trial of the efficacy of gabapentin in improving analgesia in patients undergoing hysterectomy receiving patients controlled analgesia

## Study objectives

Does gabapentin reduce morphine use and pain at rest and on movement?

## 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

Signs and Symptoms: Pain

## Interventions

1. Analgesia with gabapentin
2. Standard analgesia

## Intervention Type

Drug

## Phase

Not Applicable

## Drug/device/biological/vaccine name(s)

Gabapentin

## Primary outcome(s)

Morphine use and pain measured by visual analog score

## Key secondary outcome(s)

Not provided at time of registration

## Completion date

01/07/2007

# Eligibility

## Key inclusion criteria

Patients from Anaesthetics

## Participant type(s)

Patient

## Healthy volunteers allowed

No

## Age group

Not Specified

## Sex

Not Specified

## Key exclusion criteria

Not provided at time of registration

## Date of first enrolment

01/10/2003

## Date of final enrolment

01/07/2007

# Locations

## Countries of recruitment

United Kingdom

England

## Study participating centre

**The Middlesex Hospital**

London

United Kingdom

W1N 8AA

# Sponsor information

## Organisation

Department of Health

# Funder(s)

## Funder type

Government

## Funder Name

UCL Hospitals NHS Foundation Trust (UK)

## Funder Name

NHS R&D Support Funding

# Results and Publications

Individual participant data (IPD) sharing plan

## IPD sharing plan summary

Not provided at time of registration