

A randomised double-blind placebo-controlled trial of the efficacy of amitryptiline in post-thoracotomy chronic pain

Submission date 12/09/2003	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered
		<input type="checkbox"/> Protocol
Registration date 12/09/2003	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan
		<input type="checkbox"/> Results
Last Edited 28/11/2014	Condition category 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 M Nicholls

Contact details

Anaesthetics
Department of Anaesthetics
The Middlesex Hospital
Mortimer Street
London
United Kingdom
W1N 8AA

Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

N0263115157

Study information

Scientific Title

A randomised double-blind placebo-controlled trial of the efficacy of amitriptyline in post-thoracotomy chronic pain

Study objectives

Is amitriptyline useful as prophylaxis against post-thoracotomy pain?

Ethics approval required

Old ethics approval format

Ethics approval(s)

Not provided at time of registration

Study design

Randomised double-blind placebo-controlled trial

Primary study design

Interventional

Secondary study design

Randomised controlled trial

Study setting(s)

Hospital

Study type(s)

Treatment

Participant information sheet

Health condition(s) or problem(s) studied

Signs and Symptoms: Pain

Interventions

1. Amitriptyline
2. Placebo

Intervention Type

Drug

Drug/device/biological/vaccine name(s)

Amitriptyline

Primary outcome measure

1. Numerical verbal pain score
2. Patient pain index
3. Short McGill questionnaire

Secondary outcome measures

Not provided at time of registration

Overall study start date

24/07/2002

Completion date

01/10/2005

Eligibility

Key inclusion criteria

Not provided at time of registration

Participant type(s)

Patient

Age group

Not Specified

Sex

Not Specified

Target number of participants

60 patients from the Anaesthetics Department

Key exclusion criteria

Not provided at time of registration

Date of first enrolment

24/07/2002

Date of final enrolment

01/10/2005

Locations

Countries of recruitment

England

United Kingdom

Study participating centre

The Middlesex Hospital

London

United Kingdom

W1N 8AA

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

University College London Hospitals NHS Trust (UK)

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