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Randomised double-blind prospective controlled trial of intercostal nerve block for post-operative pain after bilateral thoracoscopic sympathectomy

Submission date 30/09/2004	**Recruitment status** No longer recruiting	[ ] Prospectively registered [ ] Protocol
Registration date 30/09/2004	**Overall study status** Completed	[ ] Statistical analysis plan [ ] Results
Last Edited 21/03/2017	**Condition category** Signs and Symptoms	[ ] Individual participant data [ ] 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
N0123138366

Study information

Scientific Title
Randomised double-blind prospective controlled trial of intercostal nerve block for post-operative pain after bilateral thoracoscopic sympathectomy

Study objectives
To assess the efficacy of thoracoscopic intercostal nerve block by laevobupivacaine in alleviating immediate postoperative pain in patients undergoing bilateral thoracoscopic sympathectomy

Ethics approval required
Old ethics approval format

Ethics approval(s)
Not provided at time of registration

Study design
Randomised double-blind prospective 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 the contact details to request a patient information sheet

Health condition(s) or problem(s) studied
Post-operative pain

Interventions
Randomised double-blind prospective controlled trial

Intervention Type
Drug

Phase
Not Applicable

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

Primary outcome measure
An assessment of the efficacy of thoracoscopic intercostal nerve block by laevobupivacaine in alleviating immediate postoperative pain in patients undergoing bilateral thoracoscopic sympathectomy

Secondary outcome measures
Not provided at time of registration

Overall study start date
01/08/2002

Completion date
31/07/2003

Eligibility

Key inclusion criteria
Patients having undergone bilateral thoracoscopic sympathectomy

Participant type(s)
Patient

Age group
Adult

Sex
Both

Target number of participants
Not provided at time of registration

Key exclusion criteria
Not provided at time of registration

Date of first enrolment
01/08/2002

Date of final enrolment
31/07/2003

Locations

Countries of recruitment
England

United Kingdom

Study participating centre
University Hospitals of Leicester
Leicester
United Kingdom
LE1 4PW

Sponsor information

Organisation
Department of Health

Sponsor details
Richmond House
79 Whitehall
London
United Kingdom
SW1A 2NL

Sponsor type
Government

Website
http://www.dh.gov.uk/Home/fs/en

Funder(s)

Funder type
Government

Funder Name
University Hospitals of Leicester 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