

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

<b>Submission date</b> 30/09/2004	<b>Recruitment status</b> No longer recruiting	<input type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
<b>Registration date</b> 30/09/2004	<b>Overall study status</b> Completed	<input type="checkbox"/> Statistical analysis plan <input type="checkbox"/> Results
<b>Last Edited</b> 21/03/2017	<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 David Raitt

### Contact details

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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