# Randomised controlled trial of port wound and intrapleural bupivacaine analgesis versus saline

Submission date	Recruitment status	<ul><li>Prospectively registered</li></ul>
29/09/2006	No longer recruiting	☐ Protocol
Registration date	Overall study status	Statistical analysis plan
29/09/2006	Completed	Results
Last Edited	Condition category	Individual participant data
08/06/2017	Surgery	Record updated in last year

# Plain English summary of protocol

Not provided at time of registration

# Contact information

# Type(s)

Scientific

#### Contact name

Mr Basil Ammori

#### Contact details

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# Additional identifiers

**EudraCT/CTIS** number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

N0453168951

# Study information

#### Scientific Title

Randomised controlled trial of port wound and intrapleural bupivacaine analgesis versus saline

## **Study objectives**

Does the administration of local anaesthetic into the port wounds and intrapleural space in patients undergoing bilateral thoracic splanchnotomy reduce postoperative pain and analgesic requirements?

# Ethics approval required

Old ethics approval format

# Ethics approval(s)

Not provided at time of registration

## Study design

Randomised double-blind 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

Surgery: Anaesthesia

#### Interventions

Administration of local anaesthetic vs no administration of local anaesthetic

#### Intervention Type

Procedure/Surgery

#### Phase

**Not Specified** 

#### Primary outcome measure

Not provided at time of registration

## Secondary outcome measures

Not provided at time of registration

# Overall study start date

01/03/2004

# Completion date

01/03/2008

# **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

40: 20 experimental group and 20 controls

## Key exclusion criteria

Not provided at time of registration

#### Date of first enrolment

01/03/2004

## Date of final enrolment

01/03/2008

# Locations

# Countries of recruitment

England

**United Kingdom** 

# Study participating centre

MRI Central Manchester & Manchester Children's University Hospitals

Manchester

**United Kingdom** 

M13 9WL

# Sponsor information

# Organisation

Record Provided by the NHSTCT Register - 2006 Update - Department of Health

## Sponsor details

Richmond House 79 Whitehall London United Kingdom SW1A 2NL +44 (0)20 7307 2622 dhmail@doh.gsi.org.uk

# Sponsor type

Government

#### Website

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

# Funder(s)

# Funder type

Hospital/treatment centre

#### Funder Name

Central Manchester and Manchester Children's University Hospitals NHS Trust (UK)

#### **Funder Name**

**Endowment fund** 

#### **Funder Name**

NHS R&D Support Funding

# **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