

Comparison of epidural and extrapleural Marcaine® for control of post-thoracotomy pain. A randomised controlled trial.

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|----------------------------------------|---------------------------------------------------|------------------------------------------------------|
| 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 24/10/2019 | Condition category Surgery | <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
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United Kingdom
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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers
N0050007015

Study information

Scientific Title

Comparison of epidural and extrapleural Marcaine® for control of post-thoracotomy pain. A randomised controlled trial.

Study objectives

Whether extrapleural analgesia is superior to epidural analgesia in:

1. Restoring pulmonary function
2. Avoiding post-operative pain
3. Inhibiting pre-operative stress response
4. Decreasing post-operative morbidity

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

Secondary study design

Randomised controlled trial

Study setting(s)

Hospital

Study type(s)

Other

Participant information sheet

Health condition(s) or problem(s) studied

Post-surgical pain

Interventions

Randomised, controlled trial. Comparison of epidural and extrapleural Marcaine®.

Intervention Type

Drug

Phase

Not Specified

Drug/device/biological/vaccine name(s)

Bupivacaine (Marcaine®)

Primary outcome measure

Not provided at time of registration

Secondary outcome measures

Not provided at time of registration

Overall study start date

01/01/1995

Completion date

31/01/1997

Eligibility

Key inclusion criteria

Post-thoracotomy patients

Participant type(s)

Patient

Age group

Not Specified

Sex

Not Specified

Target number of participants

Not provided at time of registration

Key exclusion criteria

Does not match inclusion criteria

Date of first enrolment

01/01/1995

Date of final enrolment

31/01/1997

Locations

Countries of recruitment

England

United Kingdom

Study participating centre

Thoracic Surgery
Bradford
United Kingdom
BD9 6RJ

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

Hospital/treatment centre

Funder Name

Bradford 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