

Which is better: paravertebral (PV) or interpleural (IP) bupivacaine for breast surgery?

| | | |
|--|---|--|
| 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 04/02/2014 | 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
Dr Sean Tighe

Contact details
Countess of Chester NHS Trust
Liverpool Road
Chester
United Kingdom
CH3 1UL
+44 (0)1244 365000

Additional identifiers

Protocol serial number
N0072091240

Study information

Scientific Title

Study objectives
Not provided at time of registration

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

Study type(s)

Treatment

Health condition(s) or problem(s) studied

Surgery: Breast

Interventions

Randomisation to two groups, standardised general anaesthetic, IP or PV block, recovery, test for failure, observed for pain, nausea and vomiting (N&V), patient-controlled analgesia (PCA) morphine consumption.

Intervention Type

Drug

Phase

Not Specified

Drug/device/biological/vaccine name(s)

Bupivacaine

Primary outcome(s)

1. Visual Analogue Pain Scores at rest and on movement
2. Nausea scale and vomiting episodes
3. Morphine and other analgesic consumption
4. Time to morphine demand
5. Hospital stay

Key secondary outcome(s))

Not provided at time of registration

Completion date

01/07/2003

Eligibility**Key inclusion criteria**

All patients having major breast surgery.

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Adult

Sex

Female

Key exclusion criteria

Not provided at time of registration

Date of first enrolment

01/04/2001

Date of final enrolment

01/07/2003

Locations**Countries of recruitment**

United Kingdom

England

Study participating centre

Countess of Chester NHS Trust

Chester

United Kingdom

CH3 1UL

Sponsor information**Organisation**

Department of Health (UK)

Funder(s)**Funder type**

Government

Funder Name

Countess of Chester Hospital NHS Trust (UK)

Results and Publications

Individual participant data (IPD) sharing plan

IPD sharing plan summary

Not provided at time of registration