ISRCTN45045119 https://doi.org/10.1186/ISRCTN45045119

A prospective randomised double-blinded comparison of 0.125% and 0.0625% bupivicaine for the management of postoperative pain in patients undergoing major abdominal surgery.

Submission date
12/09/2003

Registration date
12/09/2003

Last Edited
20/02/2020

Recruitment status
No longer recruiting

Overall study status
Completed

Condition category
Surgery

[_] Prospectively registered
[_] Protocol
[_] Statistical analysis plan
[X] Results
[_] Individual participant data

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

Contact name
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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

N0301111200

Study information

Scientific Title
-

Study objectives
A comparison of two epidural infusion doses for post-operative pain relief and to explore the experience of patients who have undergone epidural pain management.

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)
Treatment

Participant information sheet
Not available in web format, please use the contact details below to request a patient information sheet

Health condition(s) or problem(s) studied
Signs and Symptoms: Post operative pain

Interventions
Randomised controlled trial and telephone interviews for patient centered phase

Intervention Type
Procedure/Surgery

Phase
Not Specified

Primary outcome measure
Pain relief response to therapy

Secondary outcome measures
Not provided at time of registration

Overall study start date
31/05/2002

Completion date
30/05/2003

Eligibility

Key inclusion criteria
86 patients undergoing major abdominal surgery nursed with epidural for postoperative pain relief, randomised to two groups

Participant type(s)
Patient

Age group
Not Specified

Sex
Not Specified

Target number of participants
86

Total final enrolment
100

Key exclusion criteria
Does not meet inclusion criteria

Date of first enrolment
31/05/2002

Date of final enrolment
30/05/2003

Locations

Countries of recruitment
England

United Kingdom

Study participating centre

Blackpool, Fylde and Wyre Hospitals NHS Trust
Blackpool
United Kingdom
FY3 8NR

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
Blackpool Fylde & Wyre 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

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Other publications	Duncan F., Haigh C., Cupitt J., Vernon P., Marshall J., Nield A. . "A prospective randomized pragmatic double-blinded 0.125% and 0.0625% bupivicaine for the management of pain after operation in patients undergoing major abdominal surgery" Acute Pain, vol.7, No.2, pp.85-93 issn: 1366-0071	01/08/2005		Yes	No
Results article	results	01/08/2005		Yes	No