

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

<b>Submission date</b> 12/09/2003	<b>Recruitment status</b> No longer recruiting	<input type="checkbox"/> Prospectively registered
<b>Registration date</b> 12/09/2003	<b>Overall study status</b> Completed	<input type="checkbox"/> Protocol
<b>Last Edited</b> 20/02/2020	<b>Condition category</b> Surgery	<input type="checkbox"/> Statistical analysis plan
		<input checked="" type="checkbox"/> Results
		<input type="checkbox"/> Individual participant data

**Plain English summary of protocol**  
Not provided at time of registration

## Contact information

**Type(s)**  
Scientific

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## Additional identifiers

**Protocol serial number**  
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

## **Study type(s)**

Treatment

## **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(s)**

Pain relief response to therapy

## **Key secondary outcome(s)**

Not provided at time of registration

## **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

**Healthy volunteers allowed**

No

**Age group**

Not Specified

**Sex**

Not Specified

**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**

United Kingdom

England

**Study participating centre**

**Blackpool, Fylde and Wyre Hospitals NHS Trust**

Blackpool

United Kingdom

FY3 8NR

**Sponsor information****Organisation**

Department of Health (UK)

**Funder(s)****Funder type**

Hospital/treatment centre

## Funder Name

Blackpool Fylde & Wyre Hospitals NHS Trust (UK)

# Results and Publications

## 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?
<a href="#">Results article</a>	results	01/08/2005		Yes	No
<a href="#">Other publications</a>	Duncan F., Haigh C., Cupitt J., Vernon P., Marshall J., Nield A. . "A prospective randomized pragmatic double-blinded 0.125% and 0.0625% bupivacaine 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