

# 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
		<input type="checkbox"/> Protocol
<b>Registration date</b> 12/09/2003	<b>Overall study status</b> Completed	<input type="checkbox"/> Statistical analysis plan
		<input checked="" type="checkbox"/> Results
<b>Last Edited</b> 20/02/2020	<b>Condition category</b> Surgery	<input type="checkbox"/> Individual participant data

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

## Contact information

**Type(s)**  
Scientific

**Contact name**  
Dr Fiona Duncan

**Contact details**  
Blackpool, Fylde and Wyre Hospitals NHS Trust  
Blackpool  
United Kingdom  
FY3 8NR  
+44 (0)1253 300000  
abc@123.com

## 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?
<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
<a href="#">Results article</a>	results	01/08/2005		Yes	No