

# A prospective blinded randomised controlled trial investigating the use of patient-controlled intravenous analgesia compared with nurse-controlled anaesthesia following first time coronary artery bypass surgery.

<b>Submission date</b> 30/09/2004	<b>Recruitment status</b> No longer recruiting	<input type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
<b>Registration date</b> 30/09/2004	<b>Overall study status</b> Completed	<input type="checkbox"/> Statistical analysis plan <input type="checkbox"/> Results
<b>Last Edited</b> 18/07/2016	<b>Condition category</b> 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

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## Additional identifiers

**EudraCT/CTIS number**

**IRAS number**

**ClinicalTrials.gov number**

**Secondary identifying numbers**

N0201139523

## **Study information**

### **Scientific Title**

A prospective blinded randomised controlled trial investigating the use of patient-controlled intravenous analgesia compared with nurse-controlled anaesthesia following first time coronary artery bypass surgery.

### **Study objectives**

Is more analgesia used in patient controlled or nurse controlled regimes?

### **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 contact details to request a participant information sheet

### **Health condition(s) or problem(s) studied**

Signs and Symptoms: Post-operative pain

### **Interventions**

Patient-controlled intravenous analgesia compared with nurse-controlled anaesthesia

### **Intervention Type**

Procedure/Surgery

**Phase**

Not Applicable

**Primary outcome measure**

Not provided at time of registration

**Secondary outcome measures**

Not provided at time of registration

**Overall study start date**

01/06/2003

**Completion date**

31/05/2004

## Eligibility

**Key inclusion criteria**

Patients undergoing first time coronary artery bypass surgery

**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**

01/06/2003

**Date of final enrolment**

31/05/2004

## Locations

**Countries of recruitment**

England

United Kingdom

**Study participating centre**  
**Royal Brompton & Harefield NHS Trust**  
Harefield  
United Kingdom  
UB9 6JH

## **Sponsor information**

**Organisation**  
Department of Health

**Sponsor details**  
Richmond House  
79 Whitehall  
London  
United Kingdom  
SW1A 2NL

**Sponsor type**  
Government

**Website**  
<http://www.dh.gov.uk/Home/fs/en>

## **Funder(s)**

**Funder type**  
Government

**Funder Name**  
Royal Brompton and Harefield 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