

# 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

**Protocol serial number**  
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

## Study type(s)

Treatment

## 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(s)

Not provided at time of registration

## Key secondary outcome(s)

Not provided at time of registration

## Completion date

31/05/2004

# Eligibility

## Key inclusion criteria

Patients undergoing first time coronary artery bypass surgery

**Participant type(s)**

Patient

**Healthy volunteers allowed**

No

**Age group**

Not Specified

**Sex**

Not Specified

**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**

United Kingdom

England

**Study participating centre**

**Royal Brompton & Harefield NHS Trust**

Harefield

United Kingdom

UB9 6JH

**Sponsor information**

**Organisation**

Department of Health

**Funder(s)**

**Funder type**  
Government

**Funder Name**  
Royal Brompton and Harefield NHS Trust (UK)

## **Results and Publications**

**Individual participant data (IPD) sharing plan**

**IPD sharing plan summary**  
Not provided at time of registration