

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.

Submission date 30/09/2004	Recruitment status No longer recruiting	☐ Prospectively registered ☐ Protocol
Registration date 30/09/2004	Overall study status Completed	☐ Statistical analysis plan ☐ Results
Last Edited 18/07/2016	Condition category Surgery	☐ Individual participant data ☐ 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

Primary study design

Interventional

Study design

Randomised controlled trial

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