ISRCTN80168835 https://doi.org/10.1186/ISRCTN80168835

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

Contact name
Dr David Royston

Contact details
Anaesthetics
Royal Brompton & Harefield NHS Trust
Harefield Hospital
Hill End Road
Harefield
United Kingdom
UB9 6JH
+44 (0)1895 823737
D.Royston@rbh.nthames.nhs.uk

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