

Randomised study of neurocognitive outcome and cerebral embolic events in patients undergoing off-pump and on-pump coronary artery bypass graft surgery

Submission date 19/09/2004	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered
Registration date 22/10/2004	Overall study status Completed	<input type="checkbox"/> Protocol
Last Edited 27/11/2015	Condition category Nervous System Diseases	<input type="checkbox"/> Statistical analysis plan
		<input checked="" type="checkbox"/> Results
		<input type="checkbox"/> Individual participant data

Plain English Summary

Not provided at time of registration

Contact information

Type(s)

Scientific

Contact name

Miss Marjan Jahangiri

Contact details

Department of Cardiac Surgery
Atkinson Morley Wing
St George's Hospital & Medical School
London
United Kingdom
SW17 0QT

Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

N/A

Study information

Scientific Title

Randomised study of neurocognitive outcome and cerebral embolic events in patients undergoing off-pump and on-pump coronary artery bypass graft surgery

Study hypothesis

1. Cerebral injury, determined by neuropsychological testing, is reduced in off-pump compared with on-pump patients
2. Perioperative embolisation is reduced in off-pump, compared with on-pump, surgery
3. Any reduction in cerebral injury is mediated by a reduction in perioperative embolisation

Ethics approval required

Old ethics approval format

Ethics approval(s)

Wandsworth Local Research Ethics Committee (ref: 01.78.6, R+D Number 00.2431), in October 2001.

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

Condition

Neurocognitive dysfunction

Interventions

Patients are divided into two groups: those undergoing on-pump and those having off-pump coronary artery surgery. Comparisons between intraoperative cerebral embolic burden and postoperative neurocognitive function are made between the two groups.

Intervention Type

Other

Phase

Not Specified

Primary outcome measure

Post-operative composite neurocognitive score at six months and three years.

Secondary outcome measures

The neurocognitive score at discharge and at six weeks, and the total intra-operative microemboli count.

Overall study start date

01/08/2002

Overall study end date

01/03/2004

Eligibility

Participant inclusion criteria

Patients undergoing first time elective coronary artery bypass surgery.

Participant type(s)

Patient

Age group

Not Specified

Sex

Both

Target number of participants

Not provided at time of registration

Participant exclusion criteria

1. Previous cerebrovascular accident or Transient Ischaemic Attack (TIA)
2. Right or left internal carotid artery stenosis more than or equal to 50%
3. Previous cardiac surgery
4. Concomitant surgery, e.g. valve replacement
5. Previous psychiatric illness, e.g. depression, schizophrenia
6. Dialysis-dependent renal failure
7. Q-wave myocardial infarction in the past six weeks
8. Very poor left ventricular function (ejection fraction less than 20%)
9. Illiteracy or non-fluency in English
10. Absence of an acoustic window for transcranial Doppler ultrasound monitoring

Recruitment start date

01/08/2002

Recruitment end date

01/03/2004

Locations

Countries of recruitment

England

United Kingdom

Study participating centre

St George's Hospital

Department of Cardiac Surgery

Blackshaw Road

London

United Kingdom

SW17 0QT

Sponsor information

Organisation

St George's Hospital (UK)

Sponsor details

Blackshaw Road

London

United Kingdom

SW17 0QT

Sponsor type

Not defined

ROR

<https://ror.org/0001ke483>

Funder(s)

Funder type

Research organisation

Funder Name

St George's Hospital Cardiothoracic Research Fund

Funder Name

The Royal College of Surgeons of England

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?
Results article	results	01/03/2004		Yes	No
Results article	results	01/08/2006		Yes	No