The effect of retrograde autologous priming of cardiopulmonary bypass circuit on jugular bulb venous oxygen saturation compared with conventional priming: a prospective randomised comparative study

Submission date 12/09/2003	Recruitment status No longer recruiting	 Prospectively registered Protocol
Registration date 12/09/2003	Overall study status Completed	 Statistical analysis plan Results
Last Edited 26/02/2020	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 H Pauli

Contact details Department of Anaesthesia Leeds General Infirmary Leeds United Kingdom LS1 3EX +44 (0)113 392 6672 abc@email.com

Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers N0436117978

Study information

Scientific Title

The effect of retrograde autologous priming of cardiopulmonary bypass circuit on jugular bulb venous oxygen saturation compared with conventional priming: a prospective randomised comparative study

Study objectives

We propose to assess and compare the coupling of cerebral metabolism and cerebral blood flow in patients undergoing coronary artery bypass grafting using moderate hypothermic cardiopulmonary bypass primed with retrograde autologous or conventional priming.

Ethics approval required Old ethics approval format

Ethics approval(s) Not provided at time of registration

Study design Randomised active controlled parallel group trial

Primary study design Interventional

Secondary study design Randomised controlled trial

Study setting(s) Hospital

Study type(s) Treatment

Participant information sheet

Health condition(s) or problem(s) studied Coronary artery bypass grafting (CABG)

Interventions

Randomised controlled trial. Random allocation to: 1. Conventional priming of cardiopulmonary bypass circuit 2. Retrograde autologous priming of bypass circuit

Intervention Type Procedure/Surgery **Phase** Not Specified

Primary outcome measure To compare SjvO2 values immediately after establishing full cardiopulmonary bypass pump flow.

Secondary outcome measures Further SjvO2 values and Lactate - Oxygen Indices.

Overall study start date 31/08/2002

Completion date 31/08/2003

Eligibility

Key inclusion criteria

20 Patients scheduled to undergo first time coronary artery bypass grafting will be recruited. They will be American Society of Anaesthesiologist grade III or less.

Participant type(s) Patient

Age group Adult

Sex Both

Target number of participants

20

Key exclusion criteria

- 1. History of cerebrovascular disease
- 2. Diabetes mellitus
- 3. Body weight less than 60 kg
- 4. Haemoglobin level of less than 11g/dl
- 5. Refusal to receive blood products

Date of first enrolment

31/08/2002

Date of final enrolment 31/08/2003

Locations

Countries of recruitment

England

United Kingdom

Study participating centre Department of Anaesthesia Leeds United Kingdom LS1 3EX

Sponsor information

Organisation Department of Health (UK)

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

Sponsor type Government

Website http://www.doh.gov.uk

Funder(s)

Funder type Hospital/treatment centre

Funder Name Leeds Teaching Hospitals 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