

Low or high magnesium concentration in intermittent warm blood cardioplegia in patients undergoing coronary artery surgery: a prospective randomised study

Submission date 23/04/2002	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
Registration date 23/04/2002	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan <input checked="" type="checkbox"/> Results
Last Edited 07/06/2011	Condition category Circulatory System	<input type="checkbox"/> Individual participant data

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

PG/02/044/13713

Study information

Scientific Title

Study objectives

A comparison between two different strategies of myocardial protection.

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

Health condition(s) or problem(s) studied

Ischaemic heart disease

Interventions

Patients will be randomised to:

1. 5 mmol/l Magnesium (Mg²⁺)

2. 16 mmol/l Mg²⁺)

In the intermittent antegrade warm blood cardioplegia

Intervention Type

Drug

Phase

Not Specified

Drug/device/biological/vaccine name(s)

Magnesium

Primary outcome measure

Not provided at time of registration

Secondary outcome measures

Not provided at time of registration

Overall study start date

01/12/2001

Completion date

01/12/2004

Eligibility

Key inclusion criteria

1. A diagnosis of ischaemic heart disease
2. Undergoing primary myocardial revascularisation with the use of cardiopulmonary bypass and cardioplegic arrest

Participant type(s)

Patient

Age group

Not Specified

Sex

Not Specified

Target number of participants

400 adult patients

Key exclusion criteria

Not provided at time of registration

Date of first enrolment

01/12/2001

Date of final enrolment

01/12/2004

Locations

Countries of recruitment

England

United Kingdom

Study participating centre
Bristol Heart Institute
Bristol
United Kingdom
BS2 8HW

Sponsor information

Organisation

British Heart Foundation (UK)

Sponsor details

14 Fitzhardinge Street
London
United Kingdom
W1H 6DH
+44 (0)20 7935 0185
research@bhf.org.uk

Sponsor type

Charity

Website

<http://www.bhf.org.uk/>

ROR

<https://ror.org/02wdwnk04>

Funder(s)

Funder type

Charity

Funder Name

British Heart Foundation (UK)

Alternative Name(s)

the_bhf, The British Heart Foundation, BHF

Funding Body Type

Private sector organisation

Funding Body Subtype

Trusts, charities, foundations (both public and private)

Location

United Kingdom

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	01/09/2011		Yes	No