

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

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

**Protocol serial number**  
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

### Study type(s)

Treatment

### Health condition(s) or problem(s) studied

Ischaemic heart disease

### Interventions

Patients will be randomised to:

1. 5 mmol/l Magnesium (Mg<sup>2+</sup>)

2. 16 mmol/l Mg<sup>2+</sup>)

In the intermittent antegrade warm blood cardioplegia

### Intervention Type

Drug

### Phase

Not Specified

### Drug/device/biological/vaccine name(s)

Magnesium

### Primary outcome(s)

Not provided at time of registration

### Key secondary outcome(s)

Not provided at time of registration

### 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

**Healthy volunteers allowed**

No

**Age group**

Not Specified

**Sex**

Not Specified

**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**

United Kingdom

England

**Study participating centre****Bristol Heart Institute**

Bristol

United Kingdom

BS2 8HW

**Sponsor information****Organisation**

British Heart Foundation (UK)

ROR

https://ror.org/02wdwnk04

## Funder(s)

### Funder type

Charity

### Funder Name

British Heart Foundation (UK)

### Alternative Name(s)

The British Heart Foundation, the\_bhf, BHF

### Funding Body Type

Private sector organisation

### Funding Body Subtype

Trusts, charities, foundations (both public and private)

### Location

United Kingdom

## Results and Publications

### 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?
<a href="#">Results article</a>	results	01/09/2011		Yes	No