

Glucose-insulin-potassium (GIK) treatment in non-diabetic females undergoing cardiac surgery trial

Submission date 12/09/2003	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered
		<input type="checkbox"/> Protocol
Registration date 12/09/2003	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan
		<input type="checkbox"/> Results
Last Edited 30/04/2018	Condition category Surgery	<input type="checkbox"/> Individual participant data
		<input type="checkbox"/> Record updated in last year

Plain English summary of protocol

Not provided at time of registration

Contact information

Type(s)

Scientific

Contact name

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Contact details

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

N0016106770

Study information

Scientific Title

Glucose-insulin-potassium (GIK) treatment in non-diabetic females undergoing cardiac surgery trial

Study objectives

Will pre-operative treatment with GIK improve the mortality and morbidity in non-diabetic women undergoing cardiac revascularization surgery?

Ethics approval required

Old ethics approval format

Ethics approval(s)

Not provided at time of registration

Study design

Randomised open-label 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 the contact details below to request a patient information sheet

Health condition(s) or problem(s) studied

Cardiac surgery

Interventions

Open randomised study.

Intervention Type

Procedure/Surgery

Phase

Not Specified

Primary outcome measure

Improved mortality and morbidity

Secondary outcome measures

Not provided at time of registration

Overall study start date

26/07/2001

Completion date

25/07/2005

Eligibility

Key inclusion criteria

400 non-diabetic females undergoing cardiac surgery

Participant type(s)

Patient

Age group

Not Specified

Sex

Female

Target number of participants

400

Key exclusion criteria

Does not match inclusion criteria

Date of first enrolment

26/07/2001

Date of final enrolment

25/07/2005

Locations

Countries of recruitment

England

United Kingdom

Study participating centre

Hammersmith Hospital

London

United Kingdom
W12 0HS

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

Hammersmith Hospital 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