Investigation of the effect of GLP-1 and left ventricular function during myocardial ischaemia

| Submission date | Recruitment status No longer recruiting | Prospectively registered | | |
|-------------------|---|-----------------------------|--|--|
| 19/05/2010 | | ☐ Protocol | | |
| Registration date | Overall study status | Statistical analysis plan | | |
| 19/05/2010 | Completed | [X] Results | | |
| Last Edited | Condition category | Individual participant data | | |
| 11/08/2015 | Circulatory System | | | |

Plain English summary of protocol

Not provided at time of registration

Contact information

Type(s)

Scientific

Contact name

Dr Davis Dutka

Contact details

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

6562

Study information

Scientific Title

Investigation of the effect of GLP-1 and left ventricular function during myocardial ischaemia

Acronym

GLP-1 and left ventricular function during ischaemia

Study objectives

The hypothesis is that infusion of glucagon-like peptide-1 will protect the heart from ischaemia and improve left ventricular (LV) function during dobutamine stress echocardiography in patients with coronary artery disease

Ethics approval required

Old ethics approval format

Ethics approval(s)

MREC approved, ref: 08/H0304/68

Study design

Single-centre non-randomised interventional treatment trial

Primary study design

Interventional

Secondary study design

Non randomised study

Study setting(s)

Hospital

Study type(s)

Treatment

Participant information sheet

Health condition(s) or problem(s) studied

Topic: Cardiovascular; Subtopic: Cardiovascular (all Subtopics); Disease: Cardiovascular

Interventions

In the active DSE, patients will receive an infusion of GLP-1 intravenously at 1.2 pmol/kg/min starting 30 minutes prior to the DSE and continuing for 30 minutes into recovery. In the control scan there will be no infusion.

Intervention Type

Drug

Phase

Not Applicable

Drug/device/biological/vaccine name(s)

Glucagon-like peptide-1

Primary outcome measure

Global LV function at peak stress

Secondary outcome measures

- 1. Global LV function at 30 minutes recovery
- 2. Regional wall LV function at 30 minutes recovery
- 3. Regional wall LV function at peak stress

Overall study start date

17/06/2009

Completion date

30/07/2010

Eligibility

Key inclusion criteria

Not provided at time of registration

Participant type(s)

Patient

Age group

Not Specified

Sex

Not Specified

Target number of participants

Planned sample size: 30; UK sample size: 30

Key exclusion criteria

Not provided at time of registration

Date of first enrolment

17/06/2009

Date of final enrolment

30/07/2010

Locations

Countries of recruitment

England

United Kingdom

Study participating centre University of Cambridge Cambridge United Kingdom CB2 2QQ

Sponsor information

Organisation

Cambridge University Hospitals NHS Foundation Trust (UK)

Sponsor details

Addenbrookes Hospital Hills Road Cambridge England United Kingdom CB2 2QQ

Sponsor type

Hospital/treatment centre

Website

http://www.cuh.org.uk/

ROR

https://ror.org/04v54gj93

Funder(s)

Funder type

Research council

Funder Name

Medical Research Council (MRC) (UK)

Alternative Name(s)

Medical Research Council (United Kingdom), UK Medical Research Council, MRC

Funding Body Type

Government organisation

Funding Body Subtype

National government

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 | 01/03/2012 | | Yes | No |
| Results article | results | 08/08/2015 | | Yes | No |