Effect of glucagon-like-peptide-1 (GLP-1) on left ventricular function during percutaneous coronary intervention (PCI)

Recruitment status No longer recruiting	Prospectively registered		
	☐ Protocol		
Overall study status	Statistical analysis plan		
Completed	[X] Results		
Condition category	[] Individual participant data		
	No longer recruiting Overall study status Completed		

Plain English summary of protocol

Not provided at time of registration

Contact information

Type(s)

Scientific

Contact name

Dr Phil Read

Contact details

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

6979; G0701720

Study information

Scientific Title

Investigation of the effect of glucagon-like-peptide-1 (GLP-1) on left ventricular function during elective coronary angioplasty and stenting

Study objectives

Assessment of the cardioprotective effects of glucagon-like-peptide-1 (GLP-1) from ischaemia during coronary angioplasty.

Ethics approval required

Old ethics approval format

Ethics approval(s)

Hertfordshire Research Ethics Committee, 24/04/2009, ref: 09/H0311/17

Study design

Randomised interventional treatment 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

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

Interventions

Assessment of left ventricular function during coronary angioplasty. In the active arm patients will receive an intravenous infusion of glucagon-like peptide-1 at 1.2 pmol/kg/min which will start after the 1st balloon inflation in the coronary artery and will run until the end of the procedure. In the control arm there will be no infusion.

Intervention Type

Drug

Phase

Not Applicable

Drug/device/biological/vaccine name(s)

Glucagon-like-peptide-1

Primary outcome measure

Left ventricular function with and without GLP-1, measured during two balloon inflations in the coronary artery which are 30 minutes apart

Secondary outcome measures

- 1. Collateral flow with and without GLP-1
- 2. Troponin I at 24 hours

Measured during two balloon inflations in the coronary artery which are 30 minutes apart.

Overall study start date

22/05/2009

Completion date

01/05/2012

Eligibility

Key inclusion criteria

- 1. Age over 18 (male or female and no upper age limit)
- 2. Able to give informed consent
- 3. On the waiting list for elective PCI to a single vessel coronary stenosis
- 4. Normal left ventricular function

Participant type(s)

Patient

Age group

Adult

Lower age limit

18 Years

Sex

Both

Target number of participants

Planned Sample Size: 40

Key exclusion criteria

- 1. Atrial fibrillation
- 2. Myocardial infarction less than 3 months previously
- 3. Previous coronary artery bypass grafts
- 4. Treatment with insulin, sitagliptin, vildagliptin or exenatide

Date of first enrolment

22/05/2009

Date of final enrolment

01/05/2012

Locations

Countries of recruitment

England

United Kingdom

Study participating centre Addenbrooke's Hospital Cambridge

United Kingdom CB2 0XY

Sponsor information

Organisation

Papworth Hospital NHS Foundation Trust (UK)

Sponsor details

Papworth Everard Cambridge England United Kingdom CB3 8RE

Sponsor type

Hospital/treatment centre

Website

http://www.papworthhospital.nhs.uk/index.php

ROR

https://ror.org/01qbebb31

Funder(s)

Funder type

Research council

Funder Name

Medical Research Council (MRC) (UK) (ref: G0701720)

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	pilot study results	01/06/2011		Yes	No
Results article	results	01/02/2015		Yes	No