

Metformin against gliclazide in patients with diabetes and heart failure

Submission date 29/04/2010	Recruitment status Stopped	<input type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
Registration date 29/04/2010	Overall study status Stopped	<input type="checkbox"/> Statistical analysis plan <input type="checkbox"/> Results
Last Edited 24/04/2019	Condition category Circulatory System	<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
Mr Mike Lammiman

Contact details
Department of Cardiology
Castle Road
Cottingham
United Kingdom
HU16 5JQ

Additional identifiers

EudraCT/CTIS number
2006-002812-87

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers
4836

Study information

Scientific Title

Metformin against gliclazide in patients with diabetes and heart failure: a randomised open label study

Acronym

MAGPDF

Study objectives

Hypothesis:

In patients with symptomatic left ventricular systolic dysfunction and type 2 diabetes, biguanide therapy compared to a sulphonylurea urea (Gliclazide), will improve ventricular function and vascular endothelial function.

Summary:

There are a large number of patients with both diabetes and heart failure in the UK because diabetes causes heart failure and likewise heart failure causes diabetes. Metformin and Gliclazide are two commonly prescribed diabetic medications in these patients but there is no information on which is better for the patient. In this study we will look at the effects of these two medications on the function of the heart muscle, blood vessels and symptoms in patients with diabetes and heart failure.

Ethics approval required

Old ethics approval format

Ethics approval(s)

Hull and East Riding Research Ethics Committee (now superseded by Leeds (West) REC), 18/07/2006, ref: 06/Q1104/98

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

Follow-up:

Patients will be reviewed each month for symptom assessment using the standardised symptom questionnaire and EQ5-D. Patients will also undergo physical examination including body weight and fat composition by impedance measurement. Blood tests (non-fasting) for serum HbA1c, U&E's and blood glucose levels. Doses of the study drug (Gliclazide or Metformin) will be increased to maximum (metformin 1 g three times a day [tds] or gliclazide 160 mg twice daily [bd]) if there is inadequate diabetic control as evidenced by blood sugar readings and HbA1c levels. If this is inadequate to maintain glucose control, then agents other than a biguanide or sulphonylurea can be added to this regimen.

At 8 months all baseline tests, as described above, will be repeated and in addition the patients will undergo dobutamine stress and recovery cardiac magnetic resonance (CMR) and forearm blood flow quantification as described below. We intend to conduct dobutamine stress and recovery CMR and forearm blood flow measurements only once at 8 months (final follow-up). We could, if requested, also do these tests at baseline but this would add an extra burden to patients and extra costs but make little further contribution to addressing the primary hypothesis of differences between treatments. The study is randomised and so the two populations should be similar, which can be assessed by comparing other baseline values.

CMR Methodology:

Cardiac magnetic resonance (CMR) imaging will be performed in a 1.5T Clinical Scanner (General Electric). Heart rate and blood pressure would be monitored throughout the imaging protocol.

Cine-CMR:

Left ventricular function and volumes will be quantified at rest and after Dobutamine stress with a breath hold gated gradient echo sequence (cine-MR). Incremental doses of dobutamine, 5 - 40 mg/kg/min, will be infused intravenously. Dobutamine infusion will be continued until patients develop symptoms of myocardial ischaemia, significant arrhythmias, or achieve target heart rate. Intravenous atropine will be used if necessary to increase heart rate. Cine-MR images will be acquired at rest and at five-minute intervals for 45 minutes during recovery following Dobutamine stress.

Gadolinium contrast CMR:

First pass gadolinium enhanced imaging with a saturation recovery turboflash sequence will be done for assessment of myocardial perfusion and delayed hyperenhancement (DE-CMR) with a segmented inversion-recovery fast gradient echo sequence for assessing myocardial viability. Studies will be conducted at the same time of day, approximately 4 hours after the last intake of medication and with a similar diet in the 3 days prior to scanning as assessed by a dietary diary.

All CMR images will be sent to a Linux-based off-line workstation for analysis. The scans will be placed in random order after the identity markers have been removed. The cine and gadolinium-enhanced CMR images will be evaluated separately by the consensus of two experienced observers.

Cine images will be used to calculate LV end-diastolic volume, end-systolic volume, ejection fraction, and myocardial mass at end-diastole with the use of analytical MRI-MASS software. Epicardial and endocardial borders of the left ventricle excluding papillary muscles and trabeculations will be traced semi-automatically in contiguous short axis slices. End-systolic volumes will be adjusted for ventricular long-axis shortening, which will be accomplished by eliminating from calculation the basal slices encompassing the left atrium at end systole.

Forearm blood flow measurement:

Contrast enhanced ultrasound (CEU) imaging of the forearm muscles in short-axis will be performed with intermittent ultraharmonic imaging (Sonos 5500, Philips Ultrasound) at a transmission frequency 1.3 MHz, and a mechanical index of 1.0. For CEU lipid-shelled decafluorobutane microbubbles (DMP 115, Bistol Myers Squibb) suspended in saline (4×10^6 ml) will be infused intravenously at 1.5 to 2.5 ml/min. CEU images will be acquired at rest (CEU-R) and after an intravenous infusion of insulin (CEU-I) for 60 minutes (1 mU/kg/min and 5 mU/kg/min). Euglycaemia will be maintained with a variable rate of 20% dextrose infusion intravenously. Heart rate, blood pressure, and blood glucose levels will be monitored every 5 to 10 minutes throughout the protocol.

Digitally acquired CEU will be analysed off-line with custom written software to quantify skeletal muscle blood volume and red blood cell velocity as described previously. Skeletal muscle blood flow reserve is then given by the ratio of blood flow measured during infusion of insulin and at rest.

Total brachial artery blood flow will be quantified by 2-D and Doppler ultrasound with a 7 MHz linear-array transducer (Sonos 5500, Philips). Brachial artery blood flow will be measured by the product of vessel cross sectional area, calculated from videocaliper measurement of the diameter and the centerline time-averaged peak velocity (cm²) determined from arterial pulsed-wave Doppler, with an angle correction of 60°.

Study entry: single randomisation only

Intervention Type

Other

Phase

Phase IV

Primary outcome measure

Difference in left ventricular end-systolic volume between patients randomised to metformin compared to gliclazide at 8 months

Secondary outcome measures

Differences between the two groups in serum NT-pro BNP levels in:

1. Time to recovery of LV function after dobutamine stress
2. Insulin mediated flow reserve quantified by contrast enhanced ultrasound and markers of endothelial function
3. Average symptom score of all monthly follow-ups, which avoids undue emphasis being placed on a single time-point (patients who are withdrawn for worsening heart failure or who die will be assigned worst rank for symptom assessment at all subsequent time-points)

Overall study start date

24/10/2006

Completion date

01/09/2010

Reason abandoned (if study stopped)

Participant recruitment issue

Eligibility

Key inclusion criteria

1. Diabetes on oral hypoglycaemic agents (OHGA)
2. Heart failure
3. Male and female, lower age limit of 18 years

Participant type(s)

Patient

Age group

Adult

Lower age limit

18 Years

Sex

Both

Target number of participants

Planned Sample Size: 60

Key exclusion criteria

1. Diabetes on insulin
2. No left ventricular function (LVF)

Date of first enrolment

24/10/2006

Date of final enrolment

01/09/2010

Locations

Countries of recruitment

England

United Kingdom

Study participating centre

Department of Cardiology

Cottingham

United Kingdom

HU16 5JQ

Sponsor information

Organisation

Hull and East Yorkshire Hospitals NHS Trust (UK)

Sponsor details

Anlaby Road
Hull
England
United Kingdom
HU3 2JZ

Sponsor type

Hospital/treatment centre

Website

<http://www.hey.nhs.uk/HomeContentWithNews.aspx?PageID=1&SectionID=1>

ROR

<https://ror.org/01b11x021>

Funder(s)**Funder type**

Charity

Funder Name

British Heart Foundation (BHF) (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