

A randomised double blind placebo controlled trial of the addition of metformin to optimal treatment in patients with chronic heart failure. Assessment by regional haemodynamics neurohumoral activation and symptomatic well being

Submission date	Recruitment status	<input type="checkbox"/> Prospectively registered
12/09/2003	No longer recruiting	<input type="checkbox"/> Protocol
Registration date	Overall study status	<input type="checkbox"/> Statistical analysis plan
12/09/2003	Completed	<input type="checkbox"/> Results
Last Edited	Condition category	<input type="checkbox"/> Individual participant data
27/10/2015	Circulatory System	<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

Protocol serial number

N0192080686

Study information

Scientific Title

A randomised double blind placebo controlled trial of the addition of metformin to optimal treatment in patients with chronic heart failure. Assessment by regional haemodynamics neurohumoral activation and symptomatic well being

Study objectives

To determine whether metformin is of benefit in the treatment of patients with congestive heart failure.

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

Cardiovascular: Congestive heart failure (CHF)

Interventions

Randomised controlled trial.

Intervention Type

Drug

Phase

Not Applicable

Drug/device/biological/vaccine name(s)

Metformin

Primary outcome(s)

1. Regional haemodynamics
2. Neurohumoral
3. Symptomatic well being

Key secondary outcome(s))

Not provided at time of registration

Completion date

31/08/2004

Eligibility

Key inclusion criteria

Total number of subjects = 20

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Not Specified

Sex

Not Specified

Key exclusion criteria

Does not match inclusion criteria

Date of first enrolment

15/02/2000

Date of final enrolment

31/08/2004

Locations

Countries of recruitment

United Kingdom

England

Study participating centre

Nottingham University Hospitals NHS Trust

Nottingham

United Kingdom

NG7 2UH

Sponsor information

Organisation

Department of Health (UK)

Funder(s)

Funder type

Hospital/treatment centre

Funder Name

Nottingham University Hospitals NHS Trust (UK)

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?
Participant information sheet	Participant information sheet	11/11/2025	11/11/2025	No	Yes