

Randomised controlled trial to compare the benefits of aerobic, resistance and a combination of aerobic and resistance exercise training on stable chronic heart failure patients (Pilot Study)

Submission date 30/09/2004	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
Registration date 30/09/2004	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan <input type="checkbox"/> Results
Last Edited 19/09/2016	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

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

N0051127202

Study information

Scientific Title

Randomised controlled trial to compare the benefits of aerobic, resistance and a combination of aerobic and resistance exercise training on stable chronic heart failure patients (Pilot Study)

Study objectives

To establish what type of exercise training best improves measures of functional capacity and quality of life in patients with Chronic Heart Failure (CHF).

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

Secondary study design

Randomised controlled trial

Study setting(s)

Hospital

Study type(s)

Quality of life

Participant information sheet

Not available in web format, please use contact details to request a participant information sheet

Health condition(s) or problem(s) studied

Cardiovascular: congestive heart failure (CHF)

Interventions

Randomised controlled trial

Intervention Type

Other

Phase

Not Applicable

Primary outcome measure

Development of a combined exercise programme should be established for CHF patients

Secondary outcome measures

Not provided at time of registration

Overall study start date

01/06/2003

Completion date

30/03/2005

Eligibility

Key inclusion criteria

Thirty eligible patients with stable chronic heart failure

Participant type(s)

Patient

Age group

Adult

Sex

Both

Target number of participants

30

Key exclusion criteria

Does not match inclusion criteria

Date of first enrolment

01/06/2003

Date of final enrolment

30/03/2005

Locations

Countries of recruitment

England

United Kingdom

Study participating centre

Brighton & Sussex University Hospitals NHS Trust (RSCH)

Brighton

United Kingdom

BN2 5BE

Sponsor information

Organisation

Department of Health

Sponsor details

Richmond House

79 Whitehall

London

United Kingdom

SW1A 2NL

Sponsor type

Government

Website

<http://www.dh.gov.uk/Home/fs/en>

Funder(s)

Funder type

Hospital/treatment centre

Funder Name

Brighton and Sussex University Hospitals 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