

The optimal pace rate setting for patients with chronic heart failure and left ventricular systolic dysfunction undergoing left ventricular or multi-site pacing

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 checked="" type="checkbox"/> Results
Last Edited 20/10/2011	Condition category Circulatory System	<input type="checkbox"/> Individual participant data

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

N0084113936

Study information

Scientific Title

Study objectives

What rate should the pace-makers be set at to cope with the bradycardia which almost invariably is induced by beta blocker?

Ethics approval required

Old ethics approval format

Ethics approval(s)

Not provided at time of registration

Study design

Partially single blinded cross over design

Primary study design

Interventional

Secondary study design

Randomised controlled trial

Study setting(s)

Not specified

Study type(s)

Not Specified

Participant information sheet

Health condition(s) or problem(s) studied

Cardiovascular: Congestive heart failure (CHF)

Interventions

The patients undergo a randomised comparison of the short term effects of different pacing modes.

Intervention Type

Other

Phase

Not Specified

Primary outcome measure

Not provided at time of registration

Secondary outcome measures

Not provided at time of registration

Overall study start date

01/03/2000

Completion date

01/03/2003

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

30 patients

Key exclusion criteria

Not provided at time of registration

Date of first enrolment

01/03/2000

Date of final enrolment

01/03/2003

Locations

Countries of recruitment

England

United Kingdom

Study participating centre

Academic Cardiology Department
Cottingham
United Kingdom
HU16 5JQ

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

Government

Funder Name

The North and South Bank Research and Development Consortium (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

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Results article	results	01/10/2006		Yes	No