

Quality of life and heart rate variability of cardiac patients receiving positive psychology interventions during cardiac rehabilitation

Submission date 29/09/2006	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered
		<input type="checkbox"/> Protocol
Registration date 29/09/2006	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan
		<input type="checkbox"/> Results
Last Edited 06/11/2014	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 Richard Stantiford

Contact details

.

.

United Kingdom

.

Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

N0051177729

Study information

Scientific Title

Study objectives

Will subjects who undertake positive psychological interventions in addition to standard phase III cardiac rehabilitation demonstrate a superior outcome?

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)

Not specified

Study type(s)

Quality of life

Participant information sheet

Health condition(s) or problem(s) studied

Cardiovascular: cardiac rehabilitation therapy

Interventions

RCT: standard care vs standard care plus psychological intervention

Intervention Type

Other

Phase

Not Applicable

Primary outcome measure

Improved heart rate variability and/or improved subject well being

Secondary outcome measures

Not provided at time of registration

Overall study start date

01/03/2006

Completion date

31/07/2006

Eligibility

Key inclusion criteria

40 patients undergoing phase III cardiac rehabilitation therapy

Participant type(s)

Patient

Age group

Adult

Sex

Both

Target number of participants

40

Key exclusion criteria

Not provided at time of registration

Date of first enrolment

01/03/2006

Date of final enrolment

31/07/2006

Locations

Countries of recruitment

United Kingdom

Study participating centre

•

•

United Kingdom

•

Sponsor information

Organisation

Record Provided by the NHSTCT Register - 2006 Update - Department of Health

Sponsor details

The Department of Health, Richmond House, 79 Whitehall
London
United Kingdom
SW1A 2NL
+44 (0)20 7307 2622
dhmail@doh.gsi.org.uk

Sponsor type

Government

Website

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

Funder(s)**Funder type**

Government

Funder Name

Brighton and Sussex University Hospitals NHS Trust (UK)

Funder Name

NHS R&D Support Funding (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