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Treatment of non cardiac chest pain: a role for hypnotherapy?

Submission date 30/08/2005	**Recruitment status** No longer recruiting	[_] Prospectively registered [_] Protocol [_] Statistical analysis plan [X] Results [_] Individual participant data
Registration date 07/09/2005	**Overall study status** Completed	
Last Edited 26/05/2010	**Condition category** Signs and Symptoms	

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
RDO/22/33

Study information

Scientific Title

Study objectives
To assess whether hypnotherapy has a role in the treatment of non cardiac chest pain

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)
Treatment

Participant information sheet

Health condition(s) or problem(s) studied
Non cardiac chest pain

Interventions
Patients will be randomised to receive either:
1. 12 sessions of Hypnotherapy
or:
2. 12 sessions of supportive listening of equal duration to hypnotherapy plus placebo medication

Intervention Type
Other

Phase
Not Specified

Primary outcome measure
Subjective global assessment of relief of symptoms compared with pre-trial status.

Secondary outcome measures
To determine whether hypnotherapy improves frequency, severity and duration of non cardiac chest pain, as well as quality of life and psychological status (visual analogue scales).

Overall study start date
01/01/2002

Completion date
01/09/2005

Eligibility

Key inclusion criteria
Patients with angina-like chest pain who have been shown to have normal coronary arteries and have no other co-existent disease.

Participant type(s)
Patient

Age group
Adult

Sex
Both

Target number of participants
50 patients

Key exclusion criteria
1.Patients with proven coronary artery disease
2.Patients with oesophageal reflux disease

Date of first enrolment
01/01/2002

Date of final enrolment
01/09/2005

Locations

Countries of recruitment
England

United Kingdom

Study participating centre
Wythenshawe Hospital
Manchester
United Kingdom
M23 9LT

Sponsor information

Organisation
Department of Health (UK)

Sponsor details
Institute for Public Health Research and Policy
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Sponsor type
Hospital/treatment centre

ROR
https://ror.org/03sbpja79

Funder(s)

Funder type
Government

Funder Name
Department of Health (UK) - NHS Exec Biomedical Funding Scheme (ref: RDO/22/33)

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
Other publications	resullts	01/11/2007		Yes	No