

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

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|--|---|---|
| <b>Submission date</b><br>30/08/2005   | <b>Recruitment status</b><br>No longer recruiting | <input type="checkbox"/> Prospectively registered<br><input type="checkbox"/> Protocol            |
| <b>Registration date</b><br>07/09/2005 | <b>Overall study status</b><br>Completed          | <input type="checkbox"/> Statistical analysis plan<br><input checked="" type="checkbox"/> Results |
| <b>Last Edited</b><br>26/05/2010       | <b>Condition category</b><br>Signs and Symptoms   | <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

**Protocol serial number**  
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

**Study type(s)**

Treatment

**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(s)**

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

**Key secondary outcome(s)**

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).

**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

**Healthy volunteers allowed**

No

**Age group**

Adult

**Sex**

All

**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**

United Kingdom

England

**Study participating centre**

**Wythenshawe Hospital**

Manchester

United Kingdom

M23 9LT

## Sponsor information

**Organisation**

Department of Health (UK)

**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

**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? |
|------------------------------------|----------|--------------|------------|----------------|-----------------|
| <a href="#">Results article</a>    | results  | 01/10/2006   |            | Yes            | No              |
| <a href="#">Other publications</a> | resultts | 01/11/2007   |            | Yes            | No              |