

Non-cardiac chest pain: a randomised controlled trial of the effects of antidepressant and acid suppression treatment

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 22/05/2017	Condition category Signs and Symptoms	<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

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

Study information

Scientific Title

Non-cardiac chest pain: a randomised controlled trial of the effects of antidepressant and acid suppression treatment

Study objectives

1. To investigate the efficacy of an antidepressant drug compared to acid blockade and placebo for symptomatic relief in non-cardiac chest pain
2. To collect data in an attempt to identify factors predicting response or non-response of these patients to this treatment
3. To develop treatment strategies for this difficult group of patients
4. To correlate effects of treatment with change pain perception and in psychiatric assessment
5. To investigate the effect of treatment on quality of life

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)

Treatment

Participant information sheet

Health condition(s) or problem(s) studied

Non-cardiac chest pain

Interventions

Antidepressant drug compared to acid blockade and placebo

Intervention Type

Drug

Phase

Not Applicable

Drug/device/biological/vaccine name(s)

Antidepressant drug, acid blockade

Primary outcome measure

Not provided at time of registration

Secondary outcome measures

Not provided at time of registration

Overall study start date

01/10/2000

Completion date

31/08/2003

Eligibility

Key inclusion criteria

Patients with non-cardiac chest pain

Participant type(s)

Patient

Age group

Not Specified

Sex

Not Specified

Target number of participants

Not provided at time of registration

Key exclusion criteria

Not provided at time of registration

Date of first enrolment

01/10/2000

Date of final enrolment

31/08/2003

Locations

Countries of recruitment

England

United Kingdom

Study participating centre
Research Office
Leicester
United Kingdom
LE5 4PW

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
University Hospitals of Leicester 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