Proton pump inhibitor (PPI) treatment of gastro-oesophageal reflux (GOR) associated with non-cardiac chest pain (NCCP)

Submission date	Recruitment status No longer recruiting	Prospectively registered		
30/09/2004		☐ Protocol		
Registration date 30/09/2004	Overall study status Completed	Statistical analysis plan		
		[X] Results		
Last Edited	Condition category	[] Individual participant data		
28/07/2017	Digestive System			

Plain English summary of protocol

Not provided at time of registration

Contact information

Type(s)

Scientific

Contact name

Dr Peter Wurm

Contact details

University Hospitals of Leicester c/o Research and Development Office Leicester General Hospital NHS Trust Leicester United Kingdom LE1 4PW +44 (0)116 258 4109 nicola.turner@uhl-tr.nhs.uk

Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

Study information

Scientific Title

Proton pump inhibitor (PPI) treatment of gastro-oesophageal reflux (GOR) associated with non-cardiac chest pain (NCCP)

Study objectives

- 1. To investigate the efficacy of a proton pump inhibitor (PPI) (omeprazole 40 mg) of non-cardiac chest pain (NCCP) in patients with pH-metric evidence of abnormal gastro-oesophageal reflux (GOR)
- 2. To clarify which factor endoscopic oesophagitis, positive symptom index, positive acid perfusion test best identifies patients

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

Gastro-oesophageal reflux disease (GORD)

Interventions

Randomised controlled trial

Intervention Type

Other

Phase

Not Applicable

Primary outcome measure

Proton pump inhibitor treatment of gastro-oesophageal reflux (GOR) associated with non cardiac chest pain (NCCP)

Secondary outcome measures

Not provided at time of registration

Overall study start date

01/08/2002

Completion date

01/04/2003

Eligibility

Key inclusion criteria

Patients associated 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

01/08/2002

Date of final enrolment

01/04/2003

Locations

Countries of recruitment

England

United Kingdom

Study participating centre

University Hospitals of Leicester

Leicester United Kingdom LE1 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

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Results article	results	01/07/2010		Yes	No