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

Submission date	Recruitment status	[ ] Prospectively registered
30/09/2004	No longer recruiting	[ ] Protocol
Registration date	Overall study status	[ ] Statistical analysis plan
30/09/2004	Completed	[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