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

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

Protocol serial number
N0123138371

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

Study type(s)
Treatment

Health condition(s) or problem(s) studied
Gastro-oesophageal reflux disease (GORD)

Interventions
Randomised controlled trial

Intervention Type
Other

Phase
Not Applicable

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

Key secondary outcome(s))
Not provided at time of registration

Completion date
01/04/2003

Eligibility

Key inclusion criteria

Patients associated with non cardiac chest pain

Participant type(s)
Patient

Healthy volunteers allowed
No

Age group
Not Specified

Sex
Not Specified

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
United Kingdom

England

Study participating centre
University Hospitals of Leicester
Leicester
United Kingdom
LE1 4PW

Sponsor information

Organisation
Department of Health

Funder(s)

Funder type
Government

Funder Name
University Hospitals of Leicester NHS Trust (UK)

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?
Results article	results	01/07/2010		Yes	No