

Eradication of *Helicobacter pylori* increases ghrelin production in the gastric mucosa

Submission date
15/02/2008

Recruitment status
No longer recruiting

☐ Prospectively registered

☐ Protocol

Registration date
10/03/2008

Overall study status
Completed

☐ Statistical analysis plan

☐ Results

Last Edited
16/08/2011

Condition category
Infections and Infestations

☐ Individual participant data

☐ 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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Contact details

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Daewha-dong

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Korea, South

411-706

Additional identifiers

Protocol serial number

N/A

Study information

Scientific Title

Study objectives

To investigate whether *Helicobacter pylori* eradication influences ghrelin/obestatin production and appetite.

Ethics approval required

Old ethics approval format

Ethics approval(s)

Institutional Review Board of Inje University, Ilsan-Paik Hospital, Clinical Research Center 05-12, South Korea. Date of approval: 28/10/2005

Study design

Randomised controlled trial

Primary study design

Interventional

Study type(s)

Prevention

Health condition(s) or problem(s) studied

Helicobacter pylori infection

Interventions

The treatment group received a triple *H. pylori* eradication regimen consisting of 20 mg esomeprazole, 1,000 mg amoxicillin and 500 mg clarithromycin, twice daily for 7 days. The control group received no medication.

Intervention Type

Drug

Phase

Not Specified

Drug/device/biological/vaccine name(s)

esomeprazole, amoxicillin, and clarithromycin

Primary outcome(s)

Ghrelin mRNA levels in gastric mucosa at baseline and 5 weeks

Key secondary outcome(s)

The following were assessed at baseline and 5 weeks:

1. Obestatin and TNF- α mRNA of gastric mucosa
2. Plasma ghrelin, obestatin and ghrelin/obestatin ratio

Completion date

30/05/2006

Eligibility

Key inclusion criteria

Healthy volunteers with H. pylori infection, who saw a poster describing the study and wanted to participate.

Participant type(s)

Healthy volunteer

Healthy volunteers allowed

No

Age group

Adult

Sex

All

Key exclusion criteria

1. Age <20 or >70 years
2. Pregnancy
3. Abnormal gastric lesion, including ulcer and cancer
4. Duodenal ulcer
5. Liver disease
6. Renal impairment
7. Previous gastrointestinal surgery
8. History of H. pylori eradication
9. Drug abuse

Date of first enrolment

17/02/2006

Date of final enrolment

30/05/2006

Locations**Countries of recruitment**

Korea, South

Study participating centre

2240

Gyeonggi-do

Korea, South

411-706

Sponsor information

Organisation

Inje University, Ilsan Paik Hospital (Korea, South)

ROR

<https://ror.org/01zx5ww52>

Funder(s)**Funder type**

University/education

Funder Name

Inje University, Ilsan paik Hospital (Korea, South)

Results and Publications**Individual participant data (IPD) sharing plan****IPD sharing plan summary**

Not provided at time of registration