Eradication of Helicobacter pylori and recurrence of bleeding peptic ulcers

Submission date 10/10/2002	Recruitment status Stopped	Prospectively registeredProtocol
Registration date	Overall study status	Statistical analysis plan
10/10/2002	Stopped	Results
Last Edited	Condition category	☐ Individual participant data
22/10/2010	Digestive System	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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Additional identifiers

Protocol serial number 511003

Study information

Scientific Title

Study objectives

Not provided at time of registration

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

Recurrence of bleeding peptic ulcers

Interventions

Added as of 09/11/2009: Please note that this trial did not start as similar studies were carried out by other research groups.

Interventions:

Patients were randomised to receive:

- 1. Either a one week course of triple therapy with bismuth subcitrate, metronidazole, and tetracycline plus ranitidine, or
- 2. A six week course of ranitidine 300 mg/day.

After the ulcers healed, the antibiotic-treated patients were not given any medication whereas the ranitidine-treated patients continued to receive a maintenance dose of ranitidine 150 mg /day.

Intervention Type

Drug

Phase

Not Specified

Drug/device/biological/vaccine name(s)

Bismuth subcitrate, metronidazole, tetracycline, ranitidine

Primary outcome(s)

Not provided at time of registration

Key secondary outcome(s))

Not provided at time of registration

Completion date

01/10/2003

Reason abandoned (if study stopped)

Objectives no longer viable

Eligibility

Key inclusion criteria

- 1. Patients over the age of 16
- 2. Confirmed bleeding from either Duodenal (DU) or Gastric (GU) ulcers with or without stigmata of recent haemorrhage
- 3. A positive rapid urease test

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Adult

Sex

All

Key exclusion criteria

Not provided at time of registration

Date of first enrolment

01/10/2002

Date of final enrolment

01/10/2003

Locations

Countries of recruitment

United Kingdom

Hong Kong

Study participating centre Department of Medicine & Therapeutics

Sha Tin Hong Kong

Sponsor information

Organisation

Hong Kong Health Services Research Fund (Hong Kong)

ROR

https://ror.org/03qh32912

Funder(s)

Funder type

Research organisation

Funder Name

Hong Kong Health Services Research Fund (Hong Kong)

Results and Publications

Individual participant data (IPD) sharing plan

IPD sharing plan summary

Not provided at time of registration