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Eradication of Helicobacter pylori and recurrence of bleeding peptic ulcers

Submission date 10/10/2002	**Recruitment status** Stopped	[ ] Prospectively registered [ ] Protocol [ ] Statistical analysis plan [ ] Results [ ] Individual participant data [ ] Record updated in last year
Registration date 10/10/2002	**Overall study status** Stopped	
Last Edited 22/10/2010	**Condition category** Digestive System	

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

Contact name
Dr J J Y Sung

Contact details
Department of Medicine & Therapeutics
Prince of Wales Hospital
30 Ngan Shing Street
Sha Tin
Hong Kong
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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

Primary study design
Interventional

Study design
Randomised controlled trial

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
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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