# Eradication of Helicobacter pylori and recurrence of bleeding peptic ulcers

Submission date	Recruitment status	Prospectively registered
10/10/2002	Stopped	☐ Protocol
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

Dr J J Y Sung

#### Contact details

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

# Additional identifiers

**EudraCT/CTIS** number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers 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

#### 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

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 measure

Not provided at time of registration

#### Secondary outcome measures

Not provided at time of registration

#### Overall study start date

01/10/2002

#### 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** 

# Age group

Adult

#### Sex

Both

# Target number of participants

Not provided at time of registration

#### 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

Hong Kong

**United Kingdom** 

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)

### Sponsor details

Health Welfare and Food Bureau Government Secretariat, HKSAR 20th floor Murray Building Garden Road

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Hong Kong

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+852 2973 8288 hsrf@hwfb.gov.hk

# Sponsor type

Research organisation

#### **ROR**

https://ror.org/03qh32912

# Funder(s)

## Funder type

Research organisation

#### **Funder Name**

Hong Kong Health Services Research Fund (Hong Kong)

# **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