

# Clinical Trial Protocol

## Iranian Registry of Clinical Trials

10 Jun 2026

### Comparison of two quadruple regimen amoxicillin, bismuth subsalicylate, pantoprazole, clarythromycin and amoxicillin, bismuth subsalicylate, pantoprazole, gemifloxacin in eradication of H.pylori infection

#### Protocol summary

##### Summary

Comparison of two quadruple regimen amoxicillin, bismuth subsalicylate, pantoprazole, clarythromycin and amoxicillin, bismuth subsalicylate, pantoprazole, gemifloxacin in eradication of H.pylori infection. This study is a randomized clinical trial in new patients with H.pylori infection. Inclusion criteria: New cases of H.pylori infection; patients between 18 to 80 years old Exclusion criteria: The pregnant and breast feeding women; gastric and esophageal malignancies; pylor stenosis; liver cirrhosis; opium addicts; hypersensitivity to amoxicillin; bismuth subsalicylate; pantoprazole; clarythromycin; gemifloxacin; using cholestyramine; renal failure; history of sizzure; favism and hematologic disorders; incomplete treatment (under 80%); consent to continue treatment; history of H.pylori treatment. One hundred eighty two patients divided to two groups. One group treated with cap amoxicillin 500 miligram two numbers every 12 hours, tab bismuth subsalicylate 120 miligram two numbers every 12 hours, tab pantoprazole 20 miligram one number daily, tab clarythromycin 500 miligram one number every 12 hours and the other group treated with cap amoxicillin 500 miligram two numbers every 12 hours, tab bismuth subsalicylate 120 miligram two numbers every 12 hours, tab pantoprazole 20 miligram one number daily, tab gemifloxacin 320 miligram one number daily for 10 days period. Eight to twelve weeks after completion of treatment, all of patients assessed about eradication of H.pylori infection through urease breath test (UBT) with 14 carbon. Finally, two groups are going to compare about succeeding in eradication of H.pylori infection.

#### General information

##### Acronym

##### IRCT registration information

IRCT registration number: **IRCT201310221155N17**

Registration date: **2014-09-03, 1393/06/12**

Registration timing: **retrospective**

Last update:

Update count: **0**

##### Registration date

2014-09-03, 1393/06/12

##### Registrant information

###### Name

Farahnaz Joukar

###### Name of organization / entity

Guilan University of Medical Sciences,  
Gastrointestinal and liver disease Research Center

###### Country

Iran (Islamic Republic of)

###### Phone

+98 13 1553 5116

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##### Recruitment status

**Recruitment complete**

##### Funding source

Guilan University of Medical Sciences

##### Expected recruitment start date

2013-10-23, 1392/08/01

##### Expected recruitment end date

2014-08-23, 1393/06/01

##### Actual recruitment start date

empty

##### Actual recruitment end date

empty

##### Trial completion date

empty

##### Scientific title

Comparison of two quadruple regimen amoxicillin, bismuth subsalicylate, pantoprazole, clarythromycin and amoxicillin, bismuth subsalicylate, pantoprazole, gemifloxacin in eradication of H.pylori infection

#### Public title

The effect of gemifloxacin in treatment of H.pylori infection

#### Purpose

Treatment

#### Inclusion/Exclusion criteria

Inclusion criteria: New cases of H.pylori infection; patients between 18 to 80 years old Exclusion criteria: The pregnant and breast feeding women; gastric and esophageal malignancies; pylor stenosis; liver cirrhosis; opium addicts; hypersensitivity to amoxicillin; bismuth subsalicylate; pantoprazole; clarythromycin; gemifloxacin; using cholestyramine; renal failure; history of sizzure; favism and hematologic disorders; incomplete treatment (under 80%); consent to continue treatment; history of H.pylori treatment

#### Age

From **18 years** old to **80 years** old

#### Gender

Both

#### Phase

3

#### Groups that have been masked

*No information*

#### Sample size

Target sample size: **182**

#### Randomization (investigator's opinion)

Randomized

#### Randomization description

#### Blinding (investigator's opinion)

Not blinded

#### Blinding description

#### Placebo

Not used

#### Assignment

Parallel

#### Other design features

### Secondary Ids

empty

### Ethics committees

#### 1

##### Ethics committee

###### Name of ethics committee

Guilan University of Medical Sciences. Ethics Committee of Gastrointestinal and Liver Diseases Resea

###### Street address

Sardar Jangal Ave, Gastrointestinal and Liver Diseases Research Center, Razi Hospital, Rasht, Iran

###### City

Rasht

###### Postal code

955655-41488

#### Approval date

2010-09-23, 1389/07/01

#### Ethics committee reference number

ش-175

### Health conditions studied

#### 1

##### Description of health condition studied

Dyspepsia

##### ICD-10 code

k30

##### ICD-10 code description

Dyspepsia

### Primary outcomes

#### 1

##### Description

Helicobacter pylori eradication

##### Timepoint

12 weeks

##### Method of measurement

Ureas Breath Test with C14

### Secondary outcomes

#### 1

##### Description

Comparison of two quadruple regimen amoxicillin, bismuth subsalicylate, pantoprazole, clarythromycin and amoxicillin, bismuth subsalicylate, pantoprazole, gemifloxacin in eradication of H.pylori infection

##### Timepoint

8-12 weeks

##### Method of measurement

UBT

### Intervention groups

#### 1

##### Description

Control group take cap amoxicillin 500 miligram two numbers every 12 hours, tab bismuth subsalicylate 120 miligram two numbers every 12 hours, tab pantoprazole 20 miligram one number daily, tab clarythromycin 500 miligram one number every 12 hours for 10 days period

##### Category

Treatment - Drugs

#### 2

##### Description

Intervention group take cap amoxicillin 500 miligram two numbers every 12 hours, tab bismuth subsalicylate 120 miligram two numbers every 12 hours, tab pantoprazole

20 miligram one number daily, tab gemifloxacin 320 miligram one number daily for 10 days period

**Category**

Treatment - Drugs

**Recruitment centers**

1

**Recruitment center**

**Name of recruitment center**

Gastrointestinal and Liver Diseases Research Center

**Full name of responsible person**

Dr. Fariborz Mansour ghanei

**Street address**

Sardar Jangal Ave, Gastrointestinal and Liver Diseases Research Center, Razi Hospital, Rasht, Iran

**City**

Rasht

**Sponsors / Funding sources**

1

**Sponsor**

**Name of organization / entity**

Guilan University of Medical Sciences

**Full name of responsible person**

Dr Rasool Tabari KHomeirani

**Street address**

Sadati st, Samjoo st, Rasht,Guilan

**City**

Rasht

**Grant name**

**Grant code / Reference number**

**Is the source of funding the same sponsor organization/entity?**

Yes

**Title of funding source**

Guilan University of Medical Sciences

**Proportion provided by this source**

100

**Public or private sector**

*empty*

**Domestic or foreign origin**

*empty*

**Category of foreign source of funding**

*empty*

**Country of origin**

**Type of organization providing the funding**

*empty*

**Person responsible for general inquiries**

**Contact**

**Name of organization / entity**

Gastrointestinal and Liver Diseases Research Center

**Full name of responsible person**

Farahnaz Joukar

**Position**

Gastrointestinal and Liver Diseases Research Center

**Other areas of specialty/work**

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**Full name of responsible person**

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**Position**

Full Professor of Internal Medicine & Gastroenterology

**Other areas of specialty/work**

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**Position**

Resident of Internal Medicine

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## **Sharing plan**

**Informed Consent Form**

*empty*

**Deidentified Individual Participant Data Set (IPD)**

**Clinical Study Report**

*empty*

*empty*

**Study Protocol**

**Analytic Code**

*empty*

*empty*

**Data Dictionary**

**Statistical Analysis Plan**

*empty*