

# Clinical Trial Protocol

## Iranian Registry of Clinical Trials

29 Jun 2026

### Comparison of the efficacy of 10-day vs. 12-day concomitant therapies for first-line *Helicobacter pylori* eradication in Iran.

#### Protocol summary

##### Study aim

Evaluation and comparison of the efficacy of 10-day versus 12-day concomitant therapy as the first-line treatment for *H. pylori* eradication in Iran.

##### Design

This is a randomized single-blind clinical trial with a control group. In this study, Two hundred patients with peptic ulcer disease and *Helicobacter pylori* infection who refer to Gastroenterology clinic of Imam Khomeini hospital of Sari, will enter the study. They will be divided into 2 groups with block randomization method. The first group will receive 10-day and the second group will receive 12-day concomitant regimen. Eight weeks after therapy, *Helicobacter pylori* eradication will be assessed by Urea breath test.

##### Settings and conduct

This is a single blind study, performed on 200 patients with peptic ulcer disease and *Helicobacter pylori* infection in Gastroenterology clinic of Imam Khomeini hospital of Sari. These patients are divided into 2 groups by block randomization method and then will be treated by either 10-day or 12-day concomitant regimen.

##### Participants/Inclusion and exclusion criteria

Two hundred patients with peptic ulcer disease and *Helicobacter pylori* infection according to pathologic assessment of gastric biopsy or rapid urease test, and no history of previous treatment for *Helicobacter pylori* eradication, will enter the study.

##### Intervention groups

The first group will receive 10-day and the second group will receive 12-day concomitant regimen composed of Pantoprazole 40 mg, Amoxicillin 1000 mg, Clarithromycin 500 mg and Metronidazole 500 mg, all given twice daily.

##### Main outcome variables

*Helicobacter pylori* eradication status

#### General information

##### Reason for update

##### Acronym

##### IRCT registration information

IRCT registration number: **IRCT20170521034070N2**  
Registration date: **2019-10-28, 1398/08/06**  
Registration timing: **retrospective**

Last update: **2019-10-28, 1398/08/06**

Update count: **0**

##### Registration date

2019-10-28, 1398/08/06

##### Registrant information

##### Name

Zohreh Bari

##### Name of organization / entity

Mazandaran University of medical sciences

##### Country

Iran (Islamic Republic of)

##### Phone

+98 911 153 6575

##### Email address

zbari@mazums.ac.ir

##### Recruitment status

**Recruitment complete**

##### Funding source

##### Expected recruitment start date

2018-12-22, 1397/10/01

##### Expected recruitment end date

2019-09-22, 1398/06/31

##### Actual recruitment start date

empty

##### Actual recruitment end date

empty

##### Trial completion date

empty

##### Scientific title

Comparison of the efficacy of 10-day vs. 12-day concomitant therapies for first-line *Helicobacter pylori*

eradication in Iran.

**Public title**

Concomitant therapies for first-line Helicobacter pylori eradication

**Purpose**

Treatment

**Inclusion/Exclusion criteria****Inclusion criteria:**

Patients with peptic ulcer disease according to endoscopic findings Presence of Helicobacter pylori is proven in gastric tissue according to pathologic assessment of gastric biopsy No history of previous treatment for Helicobacter pylori

**Exclusion criteria:**

Breast feeding History of gastrointestinal surgery History of gastrointestinal malignancy using anticoagulant drugs using anti convulsion drugs pregnancy

**Age**

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

**Gender**

Both

**Phase**

3

**Groups that have been masked**

- Data analyser

**Sample size**

Target sample size: **200**

**Randomization (investigator's opinion)**

Randomized

**Randomization description**

Block randomization was used. So that patients were classified in 5-patients blocks according to their age and gender and the first five patients will enter the 10-day regimen group and the next five patients will enter the 12-day regimen group and this classification will be continued respectively until the total number of cases would be completed.

**Blinding (investigator's opinion)**

Single blinded

**Blinding description**

The statistician is blind to the study groups.

**Placebo**

Not used

**Assignment**

Parallel

**Other design features****Secondary Ids**

empty

**Ethics committees****1****Ethics committee****Name of ethics committee**

Ethics committee of Mazandaran University of Medical Sciences

**Street address**

Amir Mazandarani street

**City**

Sari

**Province**

Mazandaran

**Postal code**

4816613135

**Approval date**

2018-06-11, 1397/03/21

**Ethics committee reference number**

IR.MAZUMS.IMAMHOSPITAL.REC.1397.031

**Health conditions studied****1****Description of health condition studied**

Gastric ulcer

**ICD-10 code**

K25

**ICD-10 code description**

Gastric ulcer

**2****Description of health condition studied**

Duodenal ulcer

**ICD-10 code**

K26

**ICD-10 code description**

Duodenal ulcer

**Primary outcomes****1****Description**

Helicobacter pylori eradication

**Timepoint**

Eight weeks after receiving the drugs, Helicobacter pylori eradication will be evaluated by Urea Breath test.

**Method of measurement**

Urea Breath test.

**Secondary outcomes**

empty

**Intervention groups****1****Description**

Patients in the intervention group will receive Pantoprazole 40 mg, Amoxicillin 1g, Clarithromycin 500 mg and Metronidazole 500 mg (all given twice daily) for ten days.

**Category**

Treatment - Drugs

## 2

### Description

Patients in the control group will receive Pantoprazole 40 mg, Amoxicillin 1g, Clarithromycin 500 mg and Metronidazole 500 mg (all given twice daily) for twelve days.

### Category

Treatment - Drugs

## Recruitment centers

### 1

#### Recruitment center

##### Name of recruitment center

Imam Khomeini Hospital

##### Full name of responsible person

Zohreh Bari

##### Street address

Amir Mazandarani street

##### City

Sari

##### Province

Mazandaran

##### Postal code

4816613135

##### Phone

+98 11 3377 1760

##### Email

zohreb252@yahoo.com

## Sponsors / Funding sources

### 1

#### Sponsor

##### Name of organization / entity

Mazandaran University of Medical Sciences

##### Full name of responsible person

Dr. Majid Saeidi

##### Street address

Moallem square

##### City

Sari

##### Province

Mazandaran

##### Postal code

4817844718

##### Phone

+98 11 3448 4800

##### Email

majsaeedi@gmail.com

#### Grant name

#### Grant code / Reference number

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

Yes

#### Title of funding source

Sari University of Medical Sciences

#### Proportion provided by this source

100

#### Public or private sector

Public

#### Domestic or foreign origin

Domestic

#### Category of foreign source of funding

empty

#### Country of origin

#### Type of organization providing the funding

Academic

## Person responsible for general inquiries

### Contact

#### Name of organization / entity

Mazandaran University of Medical Sciences

#### Full name of responsible person

Zohreh Bari

#### Position

Assistant professor

#### Latest degree

Subspecialist

#### Other areas of specialty/work

Gastroenterology and hepatology

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## Person responsible for scientific inquiries

### Contact

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Zohreh Bari

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Assistant professor

#### Latest degree

Subspecialist

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## Person responsible for updating data

### Contact

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Mazandaran University of Medical Sciences

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## Sharing plan

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

Yes - There is a plan to make this available

**Study Protocol**

Yes - There is a plan to make this available

**Statistical Analysis Plan**

Yes - There is a plan to make this available

**Informed Consent Form**

Yes - There is a plan to make this available

**Clinical Study Report**

Yes - There is a plan to make this available

**Analytic Code**

Undecided - It is not yet known if there will be a plan to make this available

**Data Dictionary**

Not applicable

**Title and more details about the data/document**

The results of this study about Helicobacter pylori eradication rate will be published after deidentification of the participants.

**When the data will become available and for how long**

immediately after being published

**To whom data/document is available**

There is no limitation.

**Under which criteria data/document could be used**

There is no limitation.

**From where data/document is obtainable**

There is no limitation.

**What processes are involved for a request to access data/document**

It will be according to guidelines of the journal that will publish the article derived from the results of this study.

**Comments**