

# Clinical Trial Protocol

## Iranian Registry of Clinical Trials

20 Jun 2026

### Comparing sequential treatment with classic triple therapy for helicobacter pylori eradication

#### Protocol summary

2010-03-14, 1388/12/23

##### Summary

The primary aim of this study is to compare eradication rate and the acceptability of a new sequential treatment regimen. We designed an open-label, randomized, parallel-arm study and 130 patients with persistent dyspepsia, who referred for upper endoscopy and their rapid urease test is positive, are recruited for helicobacter pylori eradication. Patients with history of HP eradication or on PPI or recent antibiotic therapy are excluded. Patients are randomized into two groups: sequential and classic triple therapy. Triple therapy group are prescribed omeperazol 20mg BID and amoxycilin 1000mg /12h and chlarythrimicin 500mg/12h for 10 days. Sequential treatment group are prescribed amoxycilin 1000mg/12h and omeprazol 20mg/bid for first 5 days and chlarythromicin 500mg/12h and omeprazol 20mg/BID and metronidazol 500mg/12h for next 5 days. Both groups will have PPI BD for next 20 days. After 5, 10, 30 days they are contacted and asked about symptoms and side effects of medication. Information will gather based on our check list and questionnaire. All patients who are tolerated more than 80% of treatment are included for evaluation and analysis. Four to five weeks after end of antibiotics treatment they will be send for UBT and results of HP eradication and side effects will analyse.

##### Registrant information

###### Name

Azita Ganji

###### Name of organization / entity

Mashhad University of Medical Sciences

###### Country

Iran (Islamic Republic of)

###### Phone

+98 51 1882 2327

###### Email address

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##### Recruitment status

###### Recruitment complete

##### Funding source

Mashhad university of medical sciences

##### Expected recruitment start date

2010-03-14, 1388/12/23

##### Expected recruitment end date

2010-08-23, 1389/06/01

##### Actual recruitment start date

empty

##### Actual recruitment end date

empty

##### Trial completion date

empty

#### General information

##### Acronym

##### IRCT registration information

IRCT registration number: **IRCT138810152988N1**

Registration date: **2010-03-14, 1388/12/23**

Registration timing: **retrospective**

Last update:

Update count: **0**

##### Registration date

##### Scientific title

Comparing sequential treatment with classic triple therapy for helicobacter pylori eradication

##### Public title

Sequential therapy in Helicobacter pylori eradication

##### Purpose

Treatment

##### Inclusion/Exclusion criteria

Inclusion criteria: all patients with positive rapid urease test (RUT) who had indication of helicobacter eradication.

Exclusion criteria: history of HP eradication, patients on

PPI, history of allergic reaction to our medicine, history of any antibiotic in last 4 weeks, renal failure.

**Age**

No age limit

**Gender**

Both

**Phase**

N/A

**Groups that have been masked**

No information

**Sample size**

Target sample size: **130**

**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**

Ethics committee of Mashhad university of medical sciences

**Street address**

Danshgeh street

**City**

Mashhad

**Postal code****Approval date**

empty

**Ethics committee reference number**

88203

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

Eradication of Helicobacter pylori

**ICD-10 code**

K27

**ICD-10 code description**

Peptic ulcer, site unspecified

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

Eradication of Helicobacter pylori

**ICD-10 code**

K29

**ICD-10 code description**

Gastritis and duodenitis

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

helicobacter pylori eradication

**Timepoint**

45th day

**Method of measurement**

Urease btest

**Secondary outcomes****1****Description**

side effects of treatment

**Timepoint**

day 5, 10, 30

**Method of measurement**

contact by phone and if needed visit in the clinic

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

Sequential treatment: omeprazole 20mg twice daily/Amoxicillin 1gr every 12 hours for first 5 day and in next 5 days, Omeprazole 20 mg /Clarithromycin 500 mg and Metronidazole 500 mg every 12 hours and for last 20 days just omeprazole 20 mg twice daily.

**Category**

Treatment - Drugs

**2****Description**

Classic treatment (triple therapy): omeprazole 20mg, amoxicillin 1 g, clarithromycin 500mg every 12 hours for 10 days and for last 20 days just omeprazole 20 mg twice daily.

**Category**

Treatment - Drugs

**Recruitment centers****1****Recruitment center****Name of recruitment center**

Imam Reza hospital

**Full name of responsible person****Street address****City**

Mashhad

## Sponsors / Funding sources

1

### Sponsor

**Name of organization / entity**

Vice chancellor for research, Mashhad University of Medical Sciences

**Full name of responsible person**

DR Jalil Tavakol afshari

**Street address**

Mashhad university of Medical sciences, Ghoryshi building

**City**

Mashhad

**Grant name****Grant code / Reference number****Is the source of funding the same sponsor organization/entity?**

Yes

**Title of funding source**

Vice chancellor for research, Mashhad 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**

Mashhad University of Medical Sciences

**Full name of responsible person**

Azita Ganji

**Position**

Assistant professor

**Other areas of specialty/work****Street address**

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## Person responsible for updating data

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**Position**

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## Sharing plan

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

*empty*

**Study Protocol**

*empty*

**Statistical Analysis Plan**

*empty*

**Informed Consent Form**

*empty*

**Clinical Study Report**

*empty*

**Analytic Code**

*empty*

**Data Dictionary**

*empty*