

Clinical Trial Protocol

Iranian Registry of Clinical Trials

05 Jun 2026

The effect of betahistine on post-operative nausea and vomiting in patients undergoing laparoscopic cholecystectomy

Protocol summary

Registration timing: **prospective**

Study aim

The aim of this study was to evaluate the effect of betahistine in preventing nausea and vomiting after surgery of laparoscopic cholecystectomy.

Last update: **2019-01-02, 1397/10/12**

Update count: **0**

Registration date

2019-01-02, 1397/10/12

Design

70 patients randomly entered into one of two parallel groups of study or control. Study will be double blinded.

Registrant information

Name

Ebrahim Hassani

Name of organization / entity

Urmia University of Medical Sciences

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Iran (Islamic Republic of)

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Settings and conduct

This study will be performed in Urmia Imam Khomeini Hospital .90 minutes before surgery, the intervention group will receive 8 mg of oral betahistine and placebo will be given to the control group. The anesthetic method and the drugs used for anesthesia will be the same for both groups. Up to 2 hours after the operation, every 30 minutes, ask the patients about nausea and vomiting, and they will be recorded in the questionnaire. Patients and reviewers will not be aware of the type of medication and placebo.

Recruitment status

Recruitment complete

Funding source

Participants/Inclusion and exclusion criteria

patients aged 20-60 years undergoing laparoscopic cholecystectomy. Exclusion criteria include: asthma, pheochromocytoma, gastroesophageal reflux, severe obesity, BMI 35, pregnancy, breast feeding, kidney failure, history of drug addiction and alcoholism, diabetes and gastrointestinal obstruction.

Expected recruitment start date

2019-01-21, 1397/11/01

Expected recruitment end date

2019-05-21, 1398/02/31

Intervention groups

90 minutes before surgery, the intervention group will receive 8 mg of oral beta-thiostin and placebo will be given to the control group.

Actual recruitment start date

empty

Actual recruitment end date

empty

Main outcome variables

Post-operative nausea and vomiting

Trial completion date

empty

General information

Scientific title

The effect of betahistine on post-operative nausea and vomiting in patients undergoing laparoscopic cholecystectomy

Reason for update

Acronym

IRCT registration information

IRCT registration number: **IRCT20170408033280N2**

Registration date: **2019-01-02, 1397/10/12**

Public title

The effect of betahistine on post-operative nausea and

vomiting in patients undergoing laparoscopic cholecystectomy

Purpose

Prevention

Inclusion/Exclusion criteria

Inclusion criteria:

patients undergoing laparoscopic cholecystectomy from 20 to 60 years

Exclusion criteria:

Asthma Pheochromocytoma Gastroesophageal reflux severe obesity difficult airway pregnancy lactation liver and kidney diseases history of drug addiction and alcoholism neuromuscular diseases Psychiatric disorders diabetes mellitus gastrointestinal obstruction Alcoholism

Age

From **20 years** old to **60 years** old

Gender

Both

Phase

N/A

Groups that have been masked

- Participant
- Care provider
- Investigator
- Outcome assessor

Sample size

Target sample size: **70**

Randomization (investigator's opinion)

Randomized

Randomization description

Patients will be randomly assigned to one of the two groups on the basis of last digit of the file number.

Blinding (investigator's opinion)

Double blinded

Blinding description

Medication and placebo will be administered in the same volume to patients in both groups by a collaborator, and the patient, investigator and evaluator will not be aware of it.

Placebo

Used

Assignment

Parallel

Other design features

Secondary Ids

empty

Ethics committees

1

Ethics committee

Name of ethics committee

Ethics committee of Urmia University of Medical Sciences

Street address

Urmia University of Medical Sciences

City

Urmia- Resalat Boulevard-,

Province

West Azarbaijan

Postal code

5714783744

Approval date

2018-10-03, 1397/07/11

Ethics committee reference number

IR.UMSU.REC.1397.247

Health conditions studied

1

Description of health condition studied

Post -Operative Nausea and Vomiting

ICD-10 code

ICD-10 code description

Primary outcomes

1

Description

Postoperative nausea and vomiting

Timepoint

Every 30 minutes\vsakhli

Method of measurement

Questionnaire

Secondary outcomes

empty

Intervention groups

1

Description

Intervention group:Administration of 8 mg oral Beta-histeen 90 minutes before surgery

Category

Prevention

2

Description

Control group: Oral administration of the placebo 90 minutes before surgery

Category

Placebo

Recruitment centers

1

Recruitment center

Name of recruitment center

Imam Khomeini Hospital

Full name of responsible person

Ebrahim Hassani

Street address

Ershad Avenu Imam Khmeini Hospital Anesthesiology
Department

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Sponsors / Funding sources

1

Sponsor

Name of organization / entity

Oroumia University of Medical Sciences

Full name of responsible person

Professor Iraj Mohebbi

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research@umsu.ac.ir

Grant name

Grant code / Reference number

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

Yes

Title of funding source

Oroumia 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

Oroumia University of Medical Sciences

Full name of responsible person

Ebrahim Hassani

Position

Head of Department

Latest degree

Specialist

Other areas of specialty/work

Anesthesiology

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Position

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Latest degree

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Sharing plan

Deidentified Individual Participant Data Set (IPD)

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

Study Protocol

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

Statistical Analysis Plan

Not applicable

Informed Consent Form

No - There is not a plan to make this available

Clinical Study Report

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

Analytic Code

No - There is not a plan to make this available

Data Dictionary

No - There is not a plan to make this available