Clinical Trial Protocol
Iranian Registry of Clinical Trials

17 Sep 2026

A comparative study of Dexamethasone and Metoclopramide in prevention of post nausea and vomiting after hernia inguinal surgery

Protocol summary

Summary

The purpose of this study was to compare Dexamethasone and Metoclopramide in prevention of post- inguinal hernia surgery nausea and vomiting. In a randomized, double-blind clinical trial, 60 ASA class I-II patients, aged 20-40 years old and weight lower than 100kg, undergoing general anesthesia for inguinal hernia Besat hospital were recruited and assigned to receive 10mg Metoclopramide, IV or 8mg Dexamethasone, IV, 10 min before extubation and 6 hours after operation. Nausea and vomiting were measured for 1, 3, and 6 hours after the end of the operation.

General information

Acronym

IRCT registration information

IRCT registration number: **IRCT201008254505N1**
Registration date: **2010-09-11, 1389/06/20**
Registration timing: **prospective**

Last update:
Update count: **0**

Registration date

2010-09-11, 1389/06/20

Registrant information

Name

Rasool Kavyannejad

Name of organization / entity

Hamedan University of Medical Sciences

Country

Iran (Islamic Republic of)

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Recruitment status

Recruitment complete

Funding source

Kordistan Univercity of Medical Sciences

Expected recruitment start date

2010-09-16, 1389/06/25

Expected recruitment end date

2011-09-14, 1390/06/23

Actual recruitment start date

empty

Actual recruitment end date

empty

Trial completion date

empty

Scientific title

A comparative study of Dexamethasone and Metoclopramide in prevention of post nausea and vomiting after hernia inguinal surgery

Public title

Comparison of Dexamethasone and Methoclopramide on nausea and vomiting

Purpose

Treatment

Inclusion/Exclusion criteria

Inclusion criteria: inguinal hernia elective surgery, weight<100kg, age 20-40 years, no severe or chronic illness or (Asthma, Blood pressure, Lung, Renal or liver diseases), having Class ASA I-II Exclusion criteria: hypersensitivity to investigation drugs, No cooperation, Occurrence of heart and lung disorders, Smoking, opium consumption, any drug hypersensitivity

Age

From **20 years** old to **40 years** old

Gender

Both

Phase

N/A

Groups that have been masked

No information

Sample size
Target sample size: **60**
Randomization (investigator's opinion)
Randomized
Randomization description
Blinding (investigator's opinion)
Double blinded
Blinding description
Placebo
Not used
Assignment
Parallel
Other design features

Secondary Ids

empty

Ethics committees

1

Ethics committee
Name of ethics committee
Kordistan University of Medical Sciences
Street address
Pasdaran Street, Sanandaj, Kordistan
City
Sanandaj
Postal code
Approval date
2007-07-28, 1386/05/06
Ethics committee reference number
2085

Health conditions studied

1

Description of health condition studied
Nausea and vomiting
ICD-10 code
R11
ICD-10 code description

Primary outcomes

1

Description
Nausea
Timepoint
1, 3, 6 hours after operation
Method of measurement
Asking the patient and observation

2

Description
Vomiting
Timepoint
1, 3, 6 after operation
Method of measurement
observation and checklist

Secondary outcomes

empty

Intervention groups

1

Description
10mg Metoclopramide IV before extubation and 6 hours after operation
Category
Treatment - Drugs

2

Description
8mg Dexamethasone (IV) before extubation and 6 hours after operation
Category
Treatment - Drugs

Recruitment centers

1

Recruitment center
Name of recruitment center
Besat Hospital
Full name of responsible person
Street address
City
Sanandaj

Sponsors / Funding sources

1

Sponsor
Name of organization / entity
Kordistan University of Medical Sciences
Full name of responsible person
Rasool Kavyannejad
Street address
Pasdaran Street
City
Sanandaj
Grant name
Grant code / Reference number
Is the source of funding the same sponsor organization/entity?
Yes
Title of funding source
Kordistan 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

Hamedan University of Medical Sciences

Full name of responsible person

Rasool Kavyannejad

Position

MSc Student

Other areas of specialty/work

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Person responsible for updating data

Contact

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