

Clinical Trial Protocol

Iranian Registry of Clinical Trials

02 Jun 2026

Comparing the Antiemetic Effects of Ondansetron and Metoclopramide in Patients with Minor Head Trauma

Protocol summary

Summary

Nausea and vomiting are the most common complications after minor head trauma that increases the risk of intracranial pressure rising. This controlled, randomize, double blind clinical trial, will conduct in emergency department of Alzahra Hospital. The patients older than 15 years old with minor head trauma associated with nausea and vomiting will be randomly divided into 2 groups: treatment with metoclopramide (10mg/2ml, slow injection) and treatment with ondansetron (4mg/2ml, slow injection). The comparison between the 2 groups will be done regarding antiemetic efficacy and side effects.

General information

Acronym

IRCT registration information

IRCT registration number: **IRCT2015043012072N6**

Registration date: **2015-05-24, 1394/03/03**

Registration timing: **retrospective**

Last update:

Update count: **0**

Registration date

2015-05-24, 1394/03/03

Registrant information

Name

Mehrdad Esmailian

Name of organization / entity

Isfahan University of Medical Sciences

Country

Iran (Islamic Republic of)

Phone

+98 31 3629 3482

Email address

m_esmailian@med.mui.ac.ir

Recruitment status

Recruitment complete

Funding source

Isfahan University of Medical Sciences

Expected recruitment start date

2014-01-01, 1392/10/11

Expected recruitment end date

2014-06-30, 1393/04/09

Actual recruitment start date

empty

Actual recruitment end date

empty

Trial completion date

empty

Scientific title

Comparing the Antiemetic Effects of Ondansetron and Metoclopramide in Patients with Minor Head Trauma

Public title

Ondancetron Effect at Vomiting Control in Head Trauma

Purpose

Treatment

Inclusion/Exclusion criteria

Inclusion criteria : minor head trauma; nausea and vomiting; and a triage level of 3 or higher based on emergency severity score; exclusion criteria : hemodynamic instability; pregnancy/lactation; any neurologic deficit; restlessleg syndrome; alcohol usage; consumption of any antiemetic drugs during the 8 hours prior to admission; chemotherapy or radiotherapy; inability to complete and understand study explanations or outcome measures; allergy or previous adverse reactions to metoclopramide or ondansetron;

Age

From **15 years** old to **139 years** old

Gender

Both

Phase

4

Groups that have been masked

No information

Sample size

Target sample size: 120

Randomization (investigator's opinion)

Randomized

Randomization description

Blinding (investigator's opinion)

Single blinded

Blinding description

Placebo

Not used

Assignment

Parallel

Other design features

Secondary Ids

empty

Ethics committees

1

Ethics committee

Name of ethics committee

Isfahan University of Medical Sciences

Street address

Azadi Ave.

City

Isfahan

Postal code

Approval date

2013-08-06, 1392/05/15

Ethics committee reference number

65489

Health conditions studied

1

Description of health condition studied

head Injuries

ICD-10 code

S06.9

ICD-10 code description

Intracranial injury, unspecified

2

Description of health condition studied

head Injuries

ICD-10 code

S09.9

ICD-10 code description

Unspecified injury of head

Primary outcomes

1

Description

Nausea Severity

Timepoint

before drug injection and 20 minutes after drug injection

Method of measurement

Visual Analogue Scale

Secondary outcomes

1

Description

Dystonia

Timepoint

before drug injection and 20 minutes after drug injection

Method of measurement

Physical Examination

2

Description

Fatigue

Timepoint

before drug injection and 20 minutes after drug injection

Method of measurement

Physical Examination

3

Description

Headache

Timepoint

before drug injection and 20 minutes after drug injection

Method of measurement

Visual AnalogueScale

Intervention groups

1

Description

Intervention group treat with ondansetron (4mg/2ml, slow injection). 20 minutes post drug administration, nausea level will measure again. If the severity of nausea not decrease at least by 20 mm compared to the rate before the treatment intervention, a rescue dose (4mg ondansetron) would be prescribed for the patient.

Category

Treatment - Drugs

2

Description

Cotrol group treat with metoclopramide (10mg/2ml, slow injection) and treatment with ondansetron (4mg/2ml, slow injection). 20 minutes post drug administration, nausea level will measure again.

Category

Treatment - Drugs

Recruitment centers

1

Recruitment center

Name of recruitment center

Alzahra Hospital

Full name of responsible person

Mehrdad Esmailian

Street address

Sofeh Blv.

City

Isfahan

Sponsors / Funding sources

1

Sponsor

Name of organization / entity

Isfahan University of Medical Sciences

Full name of responsible person

Reza Azizkhani

Street address

Azadi Ave.

City

Isfahan

Grant name

Grant code / Reference number

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

Yes

Title of funding source

Isfahan 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

Isfahan University of Medical Sciences

Full name of responsible person

Mehrdad Esmailian

Position

Assistant Professor of Emergency Medicine

Other areas of specialty/work

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Web page address

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