

# Clinical Trial Protocol

## Iranian Registry of Clinical Trials

10 Jun 2026

### Comparison of the effect of intravenous promethazine and diazepam on the reduction of peripheral true vertigo symptom among the patient admitted to the emergency ward in Shahid Mohammadi hospital

#### Protocol summary

##### Summary

This study is a double blind clinical trial designed to compare the effect of promethazine and diazepam regarding the ability of controlling the symptom of peripheral vertigo in emergency department. Due to uncertainty of their prescribed priority in emergency reference books, the design of this study seems essential central vertigo is the main exclusion criteria. study time is considered as a 6 month interval.

#### General information

##### Acronym

##### IRCT registration information

IRCT registration number: **IRCT2017061210330N2**  
Registration date: **2017-06-26, 1396/04/05**  
Registration timing: **prospective**

Last update:

Update count: **0**

##### Registration date

2017-06-26, 1396/04/05

##### Registrant information

###### Name

Seyed Ashkan Tabibzadeh

###### Name of organization / entity

Hormozgan University of Medical Sciences

###### Country

Iran (Islamic Republic of)

###### Phone

+98 76 1333 7192

###### Email address

dr.atabibzadeh@hums.ac.ir

##### Recruitment status

**Recruitment complete**

##### Funding source

Hormozgan University of Medical Sciences

##### Expected recruitment start date

2017-07-23, 1396/05/01

##### Expected recruitment end date

2017-12-22, 1396/10/01

##### Actual recruitment start date

empty

##### Actual recruitment end date

empty

##### Trial completion date

empty

##### Scientific title

Comparison of the effect of intravenous promethazine and diazepam on the reduction of peripheral true vertigo symptom among the patient admitted to the emergency ward in Shahid Mohammadi hospital

##### Public title

Comparison of the effect of promethazine and diazepam on the treatment of peripheral vertigo

##### Purpose

Treatment

##### Inclusion/Exclusion criteria

Inclusion criteria of study :patients admitted to Shahid Mohammadi hospital with true vertigo; patients admitted to Shahid Mohammadi hospital over 18 years old and less than 70 years old;patients who do not intake anti-vertigo drugs before coming to hospital. Exclusion criteria: pregnant women; individual who suffer from know drug's allergy; people who drink alcohol; patients who feel vertigo by using drugs; patients who suffer from advanced liver disease; patients who need to follow up with more checks.

##### Age

From **18 years** old to **70 years** old

##### Gender

Both

## Phase

2-3

## Groups that have been masked

No information

## Sample size

Target sample size: **154**

## 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

Hormozgan University of Medical Sciences

##### Street address

Bandar abbas -Jomhoori eslami Blvd-The east side of the hospital Shaid Mohammadi

##### City

Bandar abbas

##### Postal code

#### Approval date

2016-12-22, 1395/10/02

#### Ethics committee reference number

IR.HUMS.REC.1395.031

## Health conditions studied

### 1

#### Description of health condition studied

Vertigo

#### ICD-10 code

R42

#### ICD-10 code description

Dizziness and giddiness

## Primary outcomes

### 1

#### Description

Severity of vertigo

#### Timepoint

At the start of study and 30 minutes after prescription

#### Method of measurement

The visual and verbal questionnaire

## Secondary outcomes

### 1

#### Description

vomiting

#### Timepoint

the start of study-30 minutes after Prescription

#### Method of measurement

the visual and verbal questionnaire

### 2

#### Description

ataxia

#### Timepoint

the start of study-30 minutes after Prescription

#### Method of measurement

the visual and verbal questionnaire

### 3

#### Description

nystagmus

#### Timepoint

the start of study-30 minutes after Prescription

#### Method of measurement

the visual and verbal questionnaire

## Intervention groups

### 1

#### Description

Diazepam is an injection of 10 mg should be dissolved in 100 ml of normal saline solution a infusion within 20 -30 minutes.

#### Category

Treatment - Drugs

### 2

#### Description

Promethazine an injection of 25 mg should be dissolved in 100 ml of normal saline solution a infusion within 20 -30 minutes.

#### Category

Treatment - Drugs

## Recruitment centers

### 1

#### Recruitment center

##### Name of recruitment center

The emergency ward in Shahid Mohammadi hospital

##### Full name of responsible person

Seyed AshkanTabibzade

##### Street address

Shaid Mohammadi hospital, Jomhoori eslami Blvd.

##### City

Bandar Abbas

## Sponsors / Funding sources

1

### Sponsor

**Name of organization / entity**

Hormozgan University of Medical Sciences

**Full name of responsible person**

Seyed Ashkan Tabibzade

**Street address**

Ostandari Blvd -Hormozgan University of Medical Sciences

**City**

Bandar Abbas

**Grant name**

**Grant code / Reference number**

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

Yes

**Title of funding source**

Hormozgan 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**

Hormozgan University of Medical Sciences

**Full name of responsible person**

Seyed Ashkan Tabibzade

**Position**

Emergency medicine specialist

**Other areas of specialty/work**

**Street address**

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dr.tabib52@gmail.com

**Web page address**

## Person responsible for scientific inquiries

### Contact

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**Other areas of specialty/work**

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## Person responsible for updating data

### Contact

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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*