

# Clinical Trial Protocol

## Iranian Registry of Clinical Trials

10 Jun 2026

### Effectiveness of mannitol on postoperative complications after impacted mandibular third molar surgery

#### Protocol summary

##### Study aim

Surgical removal of impacted mandibular third molar is one of the most common surgical procedures in the oral cavity which usually accompanies with predictable sequels such as bleeding, swelling, pain and trismus. The aim of this study is to use mannitol and review its relationship with complications such as pain, swelling and trismus

##### Design

Randomized, double-blind, placebo-uncontrolled, single-center, phase one trial

##### Settings and conduct

In this split mouth study, 30 patients referred to Hamadan dental school with the age range of 16-64 years old with impacted mandibular third molars who had bilaterally and similar impacts are to be reviewed. At least 4 weeks interval is considered between surgeries on either side to ensure recovery of the surgery. Surgery is implemented by maxillofacial surgeon but the variables were examined by student who is blind to use or non-use of mannitol and for this reason this study is blinded both on the patient and the inspector person

##### Participants/Inclusion and exclusion criteria

Inclusion criteria: Taking no medications and drugs, healthy patients with ASA 1 category, lack of allergy to local anesthesia, having bilateral mandibular third molar with similar impaction Exclusion criteria: Poor oral hygiene, hypersensitivity to mannitol, smoking and alcohol abuse, pregnant and lactating mothers

##### Intervention groups

In this study, the side of the mandible where mannitol is infused will be considered as the treatment side while the opposite side is control that is randomly selected for each patient.

##### Main outcome variables

Pain level is recorded with VAS scale, swelling is evaluated by inflexible tape and trismus is measured by assessing the interincisal distance by digital caliper on 2, 4, 6 and 8 days

#### General information

##### Reason for update

##### Acronym

##### IRCT registration information

IRCT registration number: **IRCT20160712028892N2**

Registration date: **2019-01-17, 1397/10/27**

Registration timing: **prospective**

Last update: **2019-01-17, 1397/10/27**

Update count: **0**

##### Registration date

2019-01-17, 1397/10/27

##### Registrant information

##### Name

Omid Soltaninia

##### Name of organization / entity

dental school, Hamadan university of medical sciences

##### Country

Iran (Islamic Republic of)

##### Phone

+98 38381058

##### Email address

omid.soltaninia@umsha.ac.ir

##### Recruitment status

**Recruitment complete**

##### Funding source

##### Expected recruitment start date

2019-02-01, 1397/11/12

##### Expected recruitment end date

2020-01-16, 1398/10/26

##### Actual recruitment start date

empty

##### Actual recruitment end date

empty

##### Trial completion date

empty

## Scientific title

Effectiveness of mannitol on postoperative complications after impacted mandibular third molar surgery

## Public title

Effectiveness of mannitol on postoperative complications after impacted mandibular third molar surgery

## Purpose

Prevention

## Inclusion/Exclusion criteria

### Inclusion criteria:

Taking no medications and drugs Healthy patients with ASA 1 category Lack of allergy to local anesthesia; Having bilateral mandibular third molar with similar impaction

### Exclusion criteria:

Poor oral hygiene Smoking and alcohol abuse Pregnant and lactating mothers

## Age

From **16 years** old to **64 years** old

## Gender

Both

## Phase

3

## Groups that have been masked

- Participant
- Outcome assessor

## Sample size

Target sample size: **30**

More than 1 sample in each individual

Number of samples in each individual: **2**

Side of mandible in which mannitol is applied, is considered treatment group and Contralateral side is considered control.

## Randomization (investigator's opinion)

Randomized

## Randomization description

Choosing the third molar of the mandible in either side of the patient by throwing the coin. Simple randomization or unrestricted method will be used and randomization unit is Individual

## Blinding (investigator's opinion)

Double blinded

## Blinding description

Surgery is implemented by maxillofacial surgeon but the variables were examined by student who is blind to use or non-use of mannitol and for this reason this study is blinded both on the patient and the inspector person

## Placebo

Not used

## Assignment

Parallel

## Other design features

## Secondary Ids

empty

## Ethics committees

## 1

### Ethics committee

#### Name of ethics committee

Hamadan University of Medical Sciences

#### Street address

Shahid Fahmideh Street, Hamadan City, Hamadan, Iran

#### City

Hamadan

#### Province

Hamadan

#### Postal code

6541417838741

### Approval date

2018-12-22, 1397/10/01

### Ethics committee reference number

IR.UMSHA.REC.1397.672

## Health conditions studied

## 1

### Description of health condition studied

Correlation between complications associated with impacted mandibular third molar surgery and mannitol

### ICD-10 code

### ICD-10 code description

## Primary outcomes

## 1

### Description

Pain

### Timepoint

1,2,4,6

### Method of measurement

VAS scale

## 2

### Description

Swelling

### Timepoint

2,4,6,8

### Method of measurement

Sum of 3 distances: 1- Tragus to oral commissure; 2- Lateral canthus to Mandibular angle; 3- Tragus to Pogonion

## 3

### Description

Trismus

### Timepoint

2,4,6,8

### Method of measurement

Maximum opening measured between upper and lower incisors

## Secondary outcomes

1

### Description

Patient satisfaction

### Timepoint

8

### Method of measurement

Based on planned scale

## Intervention groups

1

### Description

In this study, side of mandible in which mannitol is infused, is considered treatment group and would be selected randomly for each patient. Surgical removal of contralateral wisdom tooth would be performed at least 4 weeks after first surgery.

### Category

Treatment - Surgery

2

### Description

Contralateral side is considered control.

### Category

N/A

## Recruitment centers

1

### Recruitment center

#### Name of recruitment center

Dental School, Hamadan University of Medical Sciences

#### Full name of responsible person

Omid Soltaninia

#### Street address

Shahid Fahmideh Street, Hamadan City, Hamadan, Iran

#### City

hamadan

#### Province

Hamadan

#### Postal code

65417838741

#### Phone

+98 81 3838 1059

#### Email

dentistry@umsha.ac.ir

#### Web page address

## Sponsors / Funding sources

1

### Sponsor

### Name of organization / entity

Hamedan University of Medical Sciences

### Full name of responsible person

Omid Soltaninia

### Street address

Hamadan university of medical Sciences, Shahid Fahmideh Street, Hamadan City, Hamadan, Iran

### City

Hamadan

### Province

Hamadan

### Postal code

65417838741

### Phone

+98 81 3525 0182

### Email

ICT@umsha.ac.ir

### Grant name

### Grant code / Reference number

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

Yes

### Title of funding source

Hamedan 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

Hamedan University of Medical Sciences

#### Full name of responsible person

Omid Soltaninia

#### Position

Asistant Professor

#### Latest degree

Specialist

#### Other areas of specialty/work

Dentistry

#### Street address

Shahid Fahmideh Street, Dental School, Hamadan, Iran

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Hamadan

#### Province

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omid.soltaninia@umsha.ac.ir

## Person responsible for scientific inquiries

### Contact

**Name of organization / entity**

Hamedan University of Medical Sciences

**Full name of responsible person**

Omid Soltaninia

**Position**

Asistant Professor

**Latest degree**

Specialist

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## Person responsible for updating data

### Contact

**Name of organization / entity**

Hamedan University of Medical Sciences

**Full name of responsible person**

Omid Soltaninia

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Asistant Professor

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## Sharing plan

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

Yes - There is a plan to make this available

**Study Protocol**

Yes - There is a plan to make this available

**Statistical Analysis Plan**

Yes - There is a plan to make this available

**Informed Consent Form**

Yes - There is a plan to make this available

**Clinical Study Report**

Yes - There is a plan to make this available

**Analytic Code**

Yes - There is a plan to make this available

**Data Dictionary**

Yes - There is a plan to make this available

**Title and more details about the data/document**

All data is shared after being unidentifiable

**When the data will become available and for how long**

Starting the access period since 2020

**To whom data/document is available**

Academic researchers

**Under which criteria data/document could be used**

Use of data to advance other similar studies in this field

**From where data/document is obtainable**

contact to: [omid.soltaninia@umsha.ac.ir](mailto:omid.soltaninia@umsha.ac.ir)

**What processes are involved for a request to access data/document**

The request is checked by the compliance officer and data will be sent if the request is verified

**Comments**