

# Clinical Trial Protocol

## Iranian Registry of Clinical Trials

10 Jun 2026

### A clinical trial on the effect of ultrasound guidance on local analgesic block of the posterior tibial nerve

#### Protocol summary

##### Summary

Objective: to compare the effectiveness of ultrasound guidance in local analgesic block of the posterior tibial nerve. Design: a block-randomized, not blinded clinical trial. Study population: patients requiring posterior tibial nerve blocks. Inclusion criteria: age over 18 years, isolated trauma in posterior tibial nerve, and lack of previous ankle deformity. Exclusion criteria: multiple trauma, ankle injury, and previous history of consumption of analgesics. Sample size: 80 cases. Study intervention: the control group received the common methods (i.e., anatomical landmarks) for posterior tibial nerve block. On the other hand, the intervention group was administered posterior tibial nerve blocks using ultrasound guidance. In this method, the anesthetic drug was injected near the nerve causing pain. Study outcome: evaluation and comparison of anesthesia level and duration based on Visual Analogue Scale and Quality Block (i.e., sensory and motor block), based on cold and pinprick, 15 and 30 minutes after the intervention in both the intervention and control groups.

#### General information

##### Acronym

##### IRCT registration information

IRCT registration number: **IRCT2017041733477N1**

Registration date: **2017-06-01, 1396/03/11**

Registration timing: **registered\_while\_recruiting**

Last update:

Update count: **0**

##### Registration date

2017-06-01, 1396/03/11

##### Registrant information

##### Name

Reza Ahmadi

##### Name of organization / entity

Mashhad University of Medical Sciences

##### Country

Iran (Islamic Republic of)

##### Phone

+98 51 3841 3007

##### Email address

ebrahimimn@mums.ac.ir

##### Recruitment status

**Recruitment complete**

##### Funding source

Vice Chancellor for Research, Mashhad University of Medical Sciences

##### Expected recruitment start date

2017-05-22, 1396/03/01

##### Expected recruitment end date

2018-05-22, 1397/03/01

##### Actual recruitment start date

empty

##### Actual recruitment end date

empty

##### Trial completion date

empty

##### Scientific title

A clinical trial on the effect of ultrasound guidance on local analgesic block of the posterior tibial nerve

##### Public title

The effect of ultrasound guidance on local analgesic block

##### Purpose

Treatment

##### Inclusion/Exclusion criteria

Inclusion criteria: age over 18 years, isolated trauma in posterior tibial nerve, and lack of previous ankle deformity. Exclusion criteria: multiple trauma, ankle injury, and previous history of consumption of analgesics.

##### Age

From **18 years** old

**Gender**

Both

**Phase**

2

**Groups that have been masked**

No information

**Sample size**

Target sample size: **80**

**Randomization (investigator's opinion)**

Randomized

**Randomization description****Blinding (investigator's opinion)**

Not blinded

**Blinding description****Placebo**

Not used

**Assignment**

Parallel

**Other design features****Secondary Ids**

empty

**Ethics committees****1****Ethics committee****Name of ethics committee**

Ethics Committee of Mashhad University of Medical Sciences

**Street address**

Vice Chancellor for Research, Mashhad University of Medical Sciences, Ghoreishi building, Daneshgah Street

**City**

Mashhad

**Postal code****Approval date**

2016-03-27, 1395/01/08

**Ethics committee reference number**

IR.MUMS.fm.REC.1395.234

**Health conditions studied****1****Description of health condition studied**

Injury of nerves at ankle

**ICD-10 code**

S94

**ICD-10 code description**

Injury of nerves at ankle and foot level

**Primary outcomes****1****Description**

Pain

**Timepoint**

15 and 30 minutes after intervention

**Method of measurement**

Based on Visual Analogue Scale (VAS)

**2****Description**

Analgesic duration

**Timepoint**

After intervention

**Method of measurement**

Minutes

**Secondary outcomes****1****Description**

Quality of nerve blocks (i.e., sensory and motor block)

**Timepoint**

15 and 30 minutes after intervention

**Method of measurement**

Physical examination

**Intervention groups****1****Description**

The intervention group: Who was administered posterior tibial nerve blocks using ultrasound guidance. In this method, the anesthetic drug is injected near the nerve, causing pain.

**Category**

Treatment - Other

**2****Description**

The control group: Who received the common methods (i.e., anatomical landmarks) for posterior tibial nerve block.

**Category**

Treatment - Other

**Recruitment centers****1****Recruitment center****Name of recruitment center**

Emam Reza Hospital

**Full name of responsible person**

Reza Ahmadi

**Street address**

Imam Reza Hospital, Imam Reza Square, Ibn Sina Street

**City**

Mashhad

## Sponsors / Funding sources

1

### Sponsor

**Name of organization / entity**

Vice Chancellor for Research, Mashhad University of Medical Sciences

**Full name of responsible person**

Mohsen Tafaghodi

**Street address**

Vice Chancellor for Research, Mashhad University of Medical Sciences, Ghoreishi building, Daneshgah Street

**City**

Mashhad

**Grant name**

**Grant code / Reference number**

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

Yes

**Title of funding source**

Vice Chancellor for Research, Mashhad 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**

Mashhad University of Medical Sciences

**Full name of responsible person**

Mohsen Ebrahimi

**Position**

Emergency Medicine

**Other areas of specialty/work**

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## Person responsible for scientific

## inquiries

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## Person responsible for updating data

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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*