

# Clinical Trial Protocol

## Iranian Registry of Clinical Trials

13 Jun 2026

### A Comparison of the preventive effect of 100 mg and 200 mg oral celecoxib on shivering during and after orthopedic lower extremity surgery under the spinal anesthesia in patients and the control group

#### Protocol summary

##### Study aim

This study aimed to compare the anti-shivering effects of 100 mg and 200 mg of this drug in the control group undergoing orthopedic lower extremity surgeries under spinal anesthesia and the patients with American Society of Anesthesiologists (ASA) 1.

##### Design

Two arm parallel group randomized trial with blinded postoperative care and outcome assessment

##### Settings and conduct

In this simple randomized double blind clinical trial that was conducted at Ayatollah Kashani Hospital in Isfahan on control group, patients undergoing elective orthopedic surgery of the lower extremity under spinal anesthesia as well as patients with ASA1 were evaluated. The first group of patients received Celecoxib 100mg three hours before SA. The second group received Celecoxib 200mg three hours before SA, and the third group received placebo three hours before SA. The incidence and severity of shivering is measured 10 minutes after SA, and subsequently at 15 minutes intervals till the end of surgery in all three groups. The medications were coded to prevent any bias. The coded medications were given by the first Anesthesiologist, and the data were recorded by the second Anesthesiologist.

##### Participants/Inclusion and exclusion criteria

Participants: the patients undergoing elective orthopedic surgery of the lower extremity under spinal anesthesia  
Entry condition: patients with ASA1, age between 18 and 65  
No-entry conditions: history of hypertension; cardiovascular disease; allergy to Coxib drugs; ...

##### Intervention groups

first group is received Celecoxib 100 mg, second group Celecoxib 200 mg and third group is received placebo

##### Main outcome variables

Incidence and severity of shivering during the operation and the recovery period based on Tsai and Cha scale

#### General information

##### Reason for update

##### Acronym

spinal

##### IRCT registration information

IRCT registration number: **IRCT20101211005362N23**

Registration date: **2020-03-25, 1399/01/06**

Registration timing: **retrospective**

Last update: **2020-03-25, 1399/01/06**

Update count: **0**

##### Registration date

2020-03-25, 1399/01/06

##### Registrant information

##### Name

Mohammadreza Safavi

##### Name of organization / entity

Anesthesiology and Critical Care Research Center,  
Isfahan University of Medical Sciences, Isfahan

##### Country

Iran (Islamic Republic of)

##### Phone

+98 31 1273 2659

##### Email address

safavi@med.mui.ac.ir

##### Recruitment status

**Recruitment complete**

##### Funding source

##### Expected recruitment start date

2016-05-21, 1395/03/01

##### Expected recruitment end date

2017-05-21, 1396/02/31

##### Actual recruitment start date

2016-05-21, 1395/03/01

##### Actual recruitment end date

2017-05-21, 1396/02/31  
**Trial completion date**  
2017-05-21, 1396/02/31

**Scientific title**  
A Comparison of the preventive effect of 100 mg and 200 mg oral celecoxib on shivering during and after orthopedic lower extremity surgery under the spinal anesthesia in patients and the control group

**Public title**  
oral celecoxib on shivering during and after orthopedic lower extremity surgery

**Purpose**  
Treatment

**Inclusion/Exclusion criteria**  
**Inclusion criteria:**  
the patients with American Society of Anesthesiologists (ASA) 1 age between 18 and 65  
**Exclusion criteria:**  
History of hypertension Sever cardiovascular disease Allergy to Coxib drugs Psychological disorder based on history Temperature less than 36 or more than 38 Celsius degree Usage of alcohol and opioids Usage of vasodilators Allergy to Aspirin

**Age**  
From **18 years** old to **65 years** old

**Gender**  
Both

**Phase**  
2-3

**Groups that have been masked**

- Care provider
- Investigator
- Outcome assessor
- Data analyser

**Sample size**  
Target sample size: **105**  
Actual sample size reached: **105**

**Randomization (investigator's opinion)**  
Randomized

**Randomization description**  
Random Allocation. Numbered balls from 1 to 105 were put in a box. Then were pulled out randomly and the number was written in a group of Celecoxib 100 mg, Celecoxib 200 mg and placebo, respectively. Each number represented the order of patients entry to the hospital and had the entrance requirements and had given the consent for entering to the study.

**Blinding (investigator's opinion)**  
Double blinded

**Blinding description**  
با مارکاتین 5/0% با سوزن L3-L422 بی حس بی نخاعی در فضای Quincke spinal کیسولهای celecoxib 3 انجام گرفت. کیسولهای SA ساعت قبل از به بیمار داده شد. این کار توسط متخصص بیهوشی انجام و داده‌ها توسط متخصص بیهوشی دوم از طریق پرسشنامه جمع‌آوری و بدین طریق کورسازی صورت می‌گرفت

**Placebo**  
Used

**Assignment**  
Parallel

## Other design features

## Secondary Ids

empty

## Ethics committees

### 1

#### Ethics committee

##### Name of ethics committee

Ethics committee of Isfahan University of Medical Sciences

##### Street address

Isfahan university of medical science, Isfahan, Iran

##### City

isfahan

##### Province

Isfahan

##### Postal code

81857647852

#### Approval date

2017-05-21, 1396/02/31

#### Ethics committee reference number

IR.MUI.MED.REC.1396.260

## Health conditions studied

### 1

#### Description of health condition studied

pain

#### ICD-10 code

T88.51

#### ICD-10 code description

Hypothermia following anesthesia

## Primary outcomes

### 1

#### Description

Incidence and severity of shivering based on Tsai and Cha scale

#### Timepoint

every 15 minutes

#### Method of measurement

objective

## Secondary outcomes

empty

## Intervention groups

### 1

#### Description

First intervention group: received Celecoxib 100mg three hours before operation.

#### Category

Treatment - Drugs

## 2

### Description

Second intervention group: received Celecoxib 200mg three hours before operation.

### Category

Treatment - Drugs

## 3

### Description

Control group: received placebo three hours before operation.

### Category

Treatment - Drugs

## Recruitment centers

### 1

#### Recruitment center

##### Name of recruitment center

Kashani hospital

##### Full name of responsible person

Seyed Mohamadreza Safavi

##### Street address

Alzahra hospital, Sofeh Avenue

##### City

Isfahan

##### Province

Isfahan

##### Postal code

8178663521

##### Phone

+98 31 3620 1992

##### Email

safavi@med.mui.ac.ir

## Sponsors / Funding sources

### 1

#### Sponsor

##### Name of organization / entity

Esfahan University of Medical Sciences

##### Full name of responsible person

Shaghayegh Haghjouy Javanmard

##### Street address

Isfahan University of Medical Science, Hezarjerib Avenue

##### City

Isfahan

##### Province

Isfahan

##### Postal code

8174673461

##### Phone

+98 31 3668 8138

##### Email

research@mui.ac.ir

#### Grant name

#### Grant code / Reference number

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

Yes

#### Title of funding source

Esfahan 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

Esfahan University of Medical Sciences

##### Full name of responsible person

Seyed Mohamadreza Safavi

##### Position

Professor

##### Latest degree

Specialist

##### Other areas of specialty/work

Anesthesiology

##### Street address

Anesthesiology Office, Alzahra Hospital, Sofeh Ave

##### City

Isfahan

##### Province

Isfahan

##### Postal code

8174675731

##### Phone

+98 31 3620 1992

##### Email

safavi@med.mui.ac.ir

## Person responsible for scientific inquiries

#### Contact

##### Name of organization / entity

Esfahan University of Medical Sciences

##### Full name of responsible person

Seyed Mohamadreza Safavi

##### Position

Professor

##### Latest degree

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Anesthesiology

##### Street address

Anesthesiology Office, Alzahra Hospital, Sofeh Ave

##### City

Isfahan  
**Province**  
Isfahan  
**Postal code**  
8174675731  
**Phone**  
+98 31 3620 1992  
**Email**  
safavi@med.mui.ac.ir

8174675731  
**Phone**  
+98 31 3620 1992  
**Fax**  
**Email**  
safavi@med.mui.ac.ir  
**Web page address**

## Person responsible for updating data

### Contact

**Name of organization / entity**  
Esfahan University of Medical Sciences  
**Full name of responsible person**  
Seyed Mohamadreza Safavi  
**Position**  
Professor of Anesthesia  
**Latest degree**  
Specialist  
**Other areas of specialty/work**  
Anesthesiology  
**Street address**  
Anesthesiology Office, Alzahra Hospital, Sofeh Ave  
**City**  
Isfahan  
**Province**  
Isfahan  
**Postal code**

## Sharing plan

### Deidentified Individual Participant Data Set (IPD)

Undecided - It is not yet known if there will be a plan to make this available

### Study Protocol

Undecided - It is not yet known if there will be a plan to make this available

### Statistical Analysis Plan

Undecided - It is not yet known if there will be a plan to make this available

### Informed Consent Form

Undecided - It is not yet known if there will be a plan to make this available

### Clinical Study Report

Undecided - It is not yet known if there will be a plan to make this available

### Analytic Code

Undecided - It is not yet known if there will be a plan to make this available

### Data Dictionary

Undecided - It is not yet known if there will be a plan to make this available