

# Clinical Trial Protocol

## Iranian Registry of Clinical Trials

10 Jun 2026

### The effect of local anesthesia with lidocaine spray on the pain control of patients under radial coronary angiography

#### Protocol summary

##### Study aim

Determining the effect of local anesthesia with lidocaine spray on pain control in patients undergoing radial coronary angiography

##### Design

A three-group clinical trial with a control group, with parallel groups, randomized with the participation of 228 patients. For randomization, permutation blocks of 6 will be used.

##### Settings and conduct

The present study will be conducted as a three-group clinical trial study with the participation of 228 patients who are candidates for coronary angiography in Bushehr City Heart Hospital in 2022. Patients will be randomly divided into three intervention and control groups using permutation blocks. In the control group, only injectable lidocaine will be used to anesthetize the needle entry area. In the first intervention group, in addition to injectable lidocaine, lidocaine spray will also be used. In the second intervention group, only lidocaine spray will be used to anesthetize the needle entry area.

##### Participants/Inclusion and exclusion criteria

Inclusion criteria: Ischemic heart disease confirmed by the attending physician, Performing radial angiography for the first time, Elective patients for angiography  
Exclusion criteria: Sensitization to lidocaine skin spray, Lack of stability of vital signs during the procedure

##### Intervention groups

Intervention groups: The first intervention group (A) will receive 6 puffs of 10% lidocaine spray 5 minutes before the angiography from a distance of 5 cm around the inside of the wrist, and then they will receive injectable lidocaine and then proceed to angiography. The second intervention group (B) will receive 6 puffs of 10% lidocaine spray 5 minutes before the angiography from a distance of 5 cm from the inside of the wrist. On the contrary, the control group and the first test group will not receive injectable lidocaine.

##### Main outcome variables

Pain severity

#### General information

##### Reason for update

##### Acronym

##### IRCT registration information

IRCT registration number: **IRCT20160110025929N39**

Registration date: **2022-09-03, 1401/06/12**

Registration timing: **prospective**

Last update: **2022-09-03, 1401/06/12**

Update count: **0**

##### Registration date

2022-09-03, 1401/06/12

##### Registrant information

##### Name

Mehdi Molavi Vardanjani

##### Name of organization / entity

##### Country

Iran (Islamic Republic of)

##### Phone

+98 81 3422 5056

##### Email address

m.molavi@umsha.ac.ir

##### Recruitment status

**Recruitment complete**

##### Funding source

##### Expected recruitment start date

2022-09-21, 1401/06/30

##### Expected recruitment end date

2023-03-20, 1401/12/29

##### Actual recruitment start date

empty

##### Actual recruitment end date

empty

##### Trial completion date

empty

### Scientific title

The effect of local anesthesia with lidocaine spray on the pain control of patients under radial coronary angiography

### Public title

The effect of local anesthesia with lidocaine spray on the pain control

### Purpose

Supportive

### Inclusion/Exclusion criteria

#### Inclusion criteria:

Ischemic heart disease confirmed by the physician  
Performing radial angiography for the first time  
Elective patients for angiography

#### Exclusion criteria:

Lack of stability of vital signs before the procedure  
Sensitization to lidocaine skin spray

### Age

From **19 years** old to **59 years** old

### Gender

Both

### Phase

N/A

### Groups that have been masked

*No information*

### Sample size

Target sample size: **228**

### Randomization (investigator's opinion)

Randomized

### Randomization description

Patients will be selected through available sampling based on the inclusion criteria. The samples will be randomly assigned to three groups (two intervention groups and one control group) using block randomization method such as AABBC, ACCBAB.

### 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 Hamadan University of Medical Sciences

##### Street address

Hamadan University of Medical Sciences, Shahid Fahmideh Street

### City

Hamadan

### Province

Hamadan

### Postal code

38698-65178

### Approval date

2022-06-11, 1401/03/21

### Ethics committee reference number

IR.UMSHA.REC.1401.266

## Health conditions studied

### 1

#### Description of health condition studied

Coronary heart disease

#### ICD-10 code

I25.1

#### ICD-10 code description

Atherosclerotic heart disease of native coronary artery

## Primary outcomes

### 1

#### Description

Pain severity

#### Timepoint

Immediately after entering the needle and 30 minutes later

#### Method of measurement

Visual analog scale

## Secondary outcomes

empty

## Intervention groups

### 1

#### Description

Intervention group(A): The first intervention group (A) will receive 6 puffs of 10% lidocaine spray 5 minutes before the angiography from a distance of 5 cm around the inside of the wrist, and then they will receive injectable lidocaine and then proceed to angiography.

#### Category

N/A

### 2

#### Description

Intervention group (B): The second intervention group receives 6 puffs of 10% lidocaine spray 5 minutes before angiography from a distance of 5 cm from inside the wrist. On the contrary, the control group and the first experimental group do not receive injectable lidocaine.

#### Category

N/A

### 3

#### Description

Control group: The control group will receive the usual anesthetic procedures before angiography, which includes 0.6-1 cc of lidocaine injection of 2% in the area around the styloid bone, and then they will pierce the radial artery with a #20 needle and place a 6-French arterial sheath and they will perform vascular angiography.

#### Category

N/A

### Recruitment centers

#### 1

##### Recruitment center

###### Name of recruitment center

Boushehr heart hospital

###### Full name of responsible person

Asghar Bahrani

###### Street address

Boushehr heart hospital, Imam Khomeini Street.

###### City

Boushehr

###### Province

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###### Phone

+98 77 3333 2946

###### Email

asghar.bahrani@gmail.com

###### Web page address

### Sponsors / Funding sources

#### 1

##### Sponsor

###### Name of organization / entity

Hamedan University of Medical Sciences

###### Full name of responsible person

Reza Shokouhi

###### Street address

Hamadan University of Medical Sciences, Shahid Fahmideh Street

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###### Phone

+98 81 3838 0535

###### Email

shokoohi@yahoo.com

###### Grant name

###### Grant code / Reference number

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

No

###### Title of funding source

Deputy of research and technology

###### 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

Sajad Ebrahimi

###### Position

Studentd

###### Latest degree

Master

###### Other areas of specialty/work

Nursery

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### Person responsible for scientific inquiries

##### Contact

###### Name of organization / entity

Hamedan University of Medical Sciences

###### Full name of responsible person

Khodayar Oshvandi

###### Position

Professor

###### Latest degree

Ph.D.

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Nursery

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## Person responsible for updating data

### Contact

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**Latest degree**  
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**Other areas of specialty/work**  
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**City**  
Hamadan  
**Province**  
Hamadan  
**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