

Clinical Trial Protocol

Iranian Registry of Clinical Trials

06 Jun 2026

Study of the success rate of infraclavicular block by concurrent sonography and nerve stimulation methods

Protocol summary

Summary

The purpose of study is to compare the success rate of infraclavicular block by both sonography and nerve stimulations methods concurrently and each alone. This study is non-random, double blind and without control or placebo that is run at one center and the patients who are persenting to Shohada Ashayer hospital and are going undergo elective surgery of forearm, elbow,wrist and hand with infraclavicular nerve block And was chosen by simple non-random method that randomly divided into three groups: Sonography, nerve stimulation, sonography and nerve stimulation Inclusion criteria:patients aged 18-85 years, I and III classes of ASA(ps-1 & ps-3), consent to participate in the project. Exclusion criteria: patients with allergies to anesthesia, local infection at injection site and coagulopathy, neurological disorders in upper limb ends, recognizable psychotic disorders, cognitive disorders, drug abuse background, long- term use of epioids, severe cardiopulmonary, diabetes, recognized neuropathy, dissatisfaction to participate in the project.Sample size was 90 people. Success rate of infraclavicular block by both sonography and nerve stimulation methods concurrently and each alone. The primary outcome quick start of numbness.

General information

Acronym

IRCT registration information

IRCT registration number: **IRCT2016090716066N5**
Registration date: **2016-10-09, 1395/07/18**
Registration timing: **registered_while_recruiting**

Last update:

Update count: **0**

Registration date

2016-10-09, 1395/07/18

Registrant information

Name

Siavash Beiranvand

Name of organization / entity

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Recruitment status

Recruitment complete

Funding source

Lorestan University of Medical Sciences, Research deputy and Technology

Expected recruitment start date

2016-10-08, 1395/07/17

Expected recruitment end date

2017-05-07, 1396/02/17

Actual recruitment start date

empty

Actual recruitment end date

empty

Trial completion date

empty

Scientific title

Study of the success rate of infraclavicular block by concurrent sonography and nerve stimulation methods

Public title

The effect of sonography and nerve stimulation in Upper limbs nerver block

Purpose

Treatment

Inclusion/Exclusion criteria

Inclusion conditions: patients aged 18-85 years; I and III classes of ASA(ps-1 & ps-3); consent to participate in the project. Exclusionconditionsb:patients with allergies to

anesthesia; local infection at injection site and coagulopathy; neurological disorders in upper limb ends; recognizable psychotic disorders; cognitive disorders; drug abuse background; long-term use of opioids; severe cardiopulmonary; diabetes, recognized neuropathy; dissatisfaction to participate in the project.

Age

From **18 years** old to **85 years** old

Gender

Both

Phase

N/A

Groups that have been masked

No information

Sample size

Target sample size: **90**

Randomization (investigator's opinion)

N/A

Randomization description

Blinding (investigator's opinion)

Triple blinded

Blinding description

Placebo

Not used

Assignment

Factorial

Other design features

Secondary Ids

empty

Ethics committees

1

Ethics committee

Name of ethics committee

Lorestan University of Medical Sciences

Street address

Deputy of Research and Technology, Integrated
Pardis, Lorestan University of Medical Sciences, 5 Km
Road Khorramabad -Tehran, Khorramabad

City

Khorramabad

Postal code

Approval date

2016-05-23, 1395/03/03

Ethics committee reference number

LUMS.REC.1395.91

Health conditions studied

1

Description of health condition studied

Brachial plexus anesthesia

ICD-10 code

Y48.3

ICD-10 code description

Local anaesthetics

Primary outcomes

1

Description

numb

Timepoint

Since the injection to numbness

Method of measurement

Stopwatch

2

Description

Immobility

Timepoint

Since the injection to immobility

Method of measurement

stopwatch

Secondary outcomes

1

Description

Blood pressure values after operation

Timepoint

During operation until recovery

Method of measurement

monitoring

2

Description

Heart beat amount

Timepoint

During operation until recovery

Method of measurement

monitoring

3

Description

Intraoperative bleeding amount

Timepoint

Since the beginning to end of operation

Method of measurement

observational

4

Description

Post-operative pain amount

Timepoint

Since anesthesia to recovery

Method of measurement

Based on the VAS scale

5

Description

Extubate time

Timepoint

After operation
Method of measurement
stopwatch

6

Description
Duration of anesthesia
Timepoint
Intraoperative
Method of measurement
stopwatch

7

Description
Duration of anesthesia
Timepoint
Since the beginning of injection to recovery
Method of measurement
stopwatch

8

Description
Opening eyes
Timepoint
After operation
Method of measurement
observation

Intervention groups

1

Description
sonography on the block infraclavicular before surgery in control group
Category
Treatment - Other

2

Description
Success rate of infraclavicular block by sonography method before surgery in Intervention group
Category
Treatment - Other

3

Description
nerve stimulation in infraclavicular block before surgery in control group
Category
Treatment - Drugs

Recruitment centers

1

Recruitment center
Name of recruitment center

Shohada Ashayer Hospital Khorramabad
Full name of responsible person
dr.siavash beiranvand
Street address
Department Of Anesthesiology, Shohada Ashayer Hospital, Enghelab street, Khorramabad
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Sponsors / Funding sources

1

Sponsor
Name of organization / entity
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Grant name
Grant code / Reference number
Is the source of funding the same sponsor organization/entity?
Yes
Title of funding source
Lorestan 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

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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