

Clinical Trial Protocol

Iranian Registry of Clinical Trials

02 Jul 2026

The impact of immersion in virtual reality on the amount of anesthetic drugs used during upper limb surgeries

Protocol summary

Study aim

Evaluating the effectiveness of immersive virtual reality as an intra-operative pain management method

Design

A two-arm parallel group randomized clinical trial, non-blinded, phase 3, with a sample size of 25 patients in each group, utilizing a pool of opaque sealed envelopes for randomization.

Settings and conduct

The research will take place at Akhtar Hospital in Tehran. Patients assigned to the virtual reality group will be exposed to a tranquil virtual environment before undergoing the supraclavicular block. They will receive comprehensive explanations regarding the block procedure and reassurances that they have the option to halt the use of the virtual reality goggles at any point during the procedure if they experience any adverse effects like nausea, vomiting, or headaches. Additionally, the dosage of midazolam (0.01 mg/kg/dose) will be adjusted according to the patient's need. The experiment will utilize the Oculus Quest 2 virtual reality headset, displaying serene landscapes encompassing plains, forests, and seas, accompanied by calming music played for the patient in both the block room and operating room.

Participants/Inclusion and exclusion criteria

Inclusion criteria: Upper limb surgical operation indication, informed consent, age between 18 to 65 years. Exclusion conditions: Any uncontrolled systemic disease, history of seizures, history of migraines, inner ear disorders

Intervention groups

Virtual Reality Group: Patients undergoing upper limb surgical procedures who are immersed in a calming virtual reality environment during nerve block and surgery using a virtual reality device. Control Group: Patients undergoing upper limb surgical procedures in the usual manner for nerve block and surgery without utilizing the virtual reality device

Main outcome variables

The amount of anesthetic drugs used during the surgery

General information

Reason for update

Acronym

IRCT registration information

IRCT registration number: **IRCT20230522058251N2**

Registration date: **2024-01-18, 1402/10/28**

Registration timing: **prospective**

Last update: **2024-01-18, 1402/10/28**

Update count: **0**

Registration date

2024-01-18, 1402/10/28

Registrant information

Name

Mohammad Hossein Mahrooz

Name of organization / entity

Country

Iran (Islamic Republic of)

Phone

+98 912 130 8261

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mohamad.mahrooz@alum.sharif.edu

Recruitment status

Recruitment complete

Funding source

Expected recruitment start date

2024-01-21, 1402/11/01

Expected recruitment end date

2024-02-19, 1402/11/30

Actual recruitment start date

empty

Actual recruitment end date

empty

Trial completion date

empty

Scientific title

The impact of immersion in virtual reality on the amount of anesthetic drugs used during upper limb surgeries

Public title

Investigating the effect of immersion in virtual reality on the amount of anesthetic drugs used during upper limb surgeries

Purpose

Treatment

Inclusion/Exclusion criteria

Inclusion criteria:

Indication for upper limb surgery Informed consent to participate in the clinical trial Age between 18 to 65 years

Exclusion criteria:

Uncontrolled systemic diseases History of seizure History of migraine History of inner ear dysfunction

Age

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

Gender

Both

Phase

3

Groups that have been masked

No information

Sample size

Target sample size: **50**

Randomization (investigator's opinion)

Randomized

Randomization description

A pool of sealed, opaque envelopes containing a note with either of two phrases, 'Intervention Group' or 'Control Group,' is prepared in advance. Patients randomly chose one of these envelopes, and based on it, are allocated to either one of the two groups: intervention or control

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 Shahid Beheshti University of medical sciences

Street address

7th Floor, Bldg No.2 SBMU, Arabi Ave, Daneshjoo Blvd, Velenjak, Tehran, Iran

City

Tehran

Province

Tehran

Postal code

1985717443

Approval date

2023-10-28, 1402/08/06

Ethics committee reference number

IR.SBMU.MSP.REC.1402.394

Health conditions studied

1

Description of health condition studied

Fracture of lower end of radius

ICD-10 code

S52.5

ICD-10 code description

Fracture of lower end of radius

2

Description of health condition studied

Carpal tunnel syndrome

ICD-10 code

G56.0

ICD-10 code description

Carpal tunnel syndrome

Primary outcomes

1

Description

The amount of anesthetic drug administered to the patient

Timepoint

before nerve block and during surgery

Method of measurement

The amount of Midazolam or Fentanyl injected to the patient as recorded in the surgery report note

Secondary outcomes

empty

Intervention groups

1

Description

Intervention group: Candidates for upper limb surgery who, before the start of supraclavicular block and during the operation, are placed under virtual reality immersion and receive fentanyl and midazolam at the discretion of the anesthesiologist

Category

Treatment - Devices

2

Description

Control group: Candidates for upper limb surgery who, during supraclavicular block and during the surgery, receive fentanyl and midazolam according to the standard protocol and under the supervision of the anesthesiologist

Category

Treatment - Drugs

Recruitment centers

1

Recruitment center

Name of recruitment center

بیمارستان ارتوپدی اختر

Full name of responsible person

Amir Bisadi

Street address

تهران - خیابان شریعتی - روبرو مترو قیطریه - خیابان شریفی
منش - بن بست آذر - بیمارستان اختر

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Tehran

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Postal code

1964714953

Phone

+98 21 2200 1072

Email

akhtarhospital@sbmu.ac.ir

Web page address

Sponsors / Funding sources

1

Sponsor

Name of organization / entity

Shahid Beheshti University of Medical Sciences

Full name of responsible person

Reza Minaei-Noshahr

Street address

Akhtar hospital, Azar dead end, Sharifimanesh st,
Shariati st, Tehran

City

Tehran

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Tehran

Postal code

1964714953

Phone

+98 21 2200 1072

Email

akhtarhospital@sbmu.ac.ir

Grant name

Grant code / Reference number

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

No

Title of funding source

Reza Minaei-Noshahr

Proportion provided by this source

100

Public or private sector

Private

Domestic or foreign origin

Domestic

Category of foreign source of funding

empty

Country of origin

Type of organization providing the funding

Persons

Person responsible for general inquiries

Contact

Name of organization / entity

Shahid Beheshti University of Medical Sciences

Full name of responsible person

Mohammad Hossein Mahrooz

Position

Medical Intern

Latest degree

Master

Other areas of specialty/work

Orthopedics

Street address

Akhtar hospital, Azar dead end, Sharifimanesh st,
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Person responsible for scientific inquiries

Contact

Name of organization / entity

Shahid Beheshti University of Medical Sciences

Full name of responsible person

Amir Bisadi

Position

Assistant Professor of Orthopaedic hand surgery

Latest degree

Subspecialist

Other areas of specialty/work

Orthopedics

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Person responsible for updating data

Contact

Name of organization / entity
Shahid Beheshti University of Medical Sciences
Full name of responsible person
Qumars Kasnavi
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Postal code
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Phone
+98 21 2200 1072
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akhtarhospital@sbm.ac.ir

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

Not applicable

Data Dictionary

Not applicable

Title and more details about the data/document

The data in this study include patients' age, gender, type of surgery, duration of the procedure, and the type and amount of anesthetic drug administered. All data will be anonymized to ensure patient confidentiality

When the data will become available and for how long

Data will be available as soon as the article is published

To whom data/document is available

Researchers in every university, or other scientific research institutes

Under which criteria data/document could be used

The data will be anonymized to ensure patient confidentiality

From where data/document is obtainable

To access data please contact the corresponding author using the contact information included in the published article

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

The request should be formally submitted in a formal letterhead from the university or research institute as the requesting party for the corresponding author

Comments