

Clinical Trial Protocol

Iranian Registry of Clinical Trials

26 Jun 2026

Comparison of the effect of perioperative intravenous ketamine with fentanyl in control of acute pain after inguinal hernia repair surgery

Protocol summary

Study aim

Comparison of the effect of perioperative intravenous ketamine with fentanyl in control of acute pain after inguinal hernia repair surgery

Design

A randomized, double-blind clinical trial with a parallel groups design; phase 1-2; on 60 patients; divided into two groups of ketamine and fentanyl (control) by random software allocation.

Settings and conduct

This study will be conducted in urmia Imam Khomeini Medical Center. After entering the study, the participants will be randomly assigned to one of two ketamine and fentanyl (control) groups with 30 participants in each group. The patients and the nurse who will take care of them during the procedure will be blinded.

Participants/Inclusion and exclusion criteria

Inclusion criteria: 30-60 aged adults with inguinal hernia, Absence of contraindications for ketamine and opioids use, patients with American Society of Anesthesiologist (ASA) physical status I,II Exclusion criteria: History of allergic reactions to the drug under study, History of underlying diseases (diabetes, peptic ulcer, gastritis, coronary artery disease, renal failure), Any known psychiatric disorder, Patients with incarcerated hernia, patients with severe infection

Intervention groups

Intervention group: In ketamine group, Induction of anesthesia is done with 1 to 4 mg/kg of intravenous ketamine, and then in maintenance phase, intravenous perfusion of ketamine at a rate of 2 to 10 µg/kg/min will be prescribed. Control group: In fentanyl group, induction of anesthesia is performed with 1-2 µg/kg fentanyl as a routine anesthetic drug, and then maintenance phase, continuous infusion of fentanyl at a rate of 1-2 µg/kg/hr will be prescribed.

Main outcome variables

Postoperative pain measurement at 60, 90 minutes and 24 hours after surgery with numeric rating (NRS) scale

from 0 to 10

General information

Reason for update

Acronym

IRCT registration information

IRCT registration number: **IRCT20221126056613N5**

Registration date: **2023-08-27, 1402/06/05**

Registration timing: **registered_while_recruiting**

Last update: **2023-08-27, 1402/06/05**

Update count: **0**

Registration date

2023-08-27, 1402/06/05

Registrant information

Name

aliakbar nasiri

Name of organization / entity

Country

Iran (Islamic Republic of)

Phone

+98 44 3346 9931

Email address

nasiriali7@gmail.com

Recruitment status

Recruitment complete

Funding source

Expected recruitment start date

2023-04-21, 1402/02/01

Expected recruitment end date

2023-09-22, 1402/06/31

Actual recruitment start date

empty

Actual recruitment end date

empty

Trial completion date

empty

Scientific title

Comparison of the effect of perioperative intravenous ketamine with fentanyl in control of acute pain after inguinal hernia repair surgery

Public title

Comparison of the effect of ketamine with fentanyl in control of acute pain after surgery

Purpose

Treatment

Inclusion/Exclusion criteria

Inclusion criteria:

30-60 aged adults with inguinal hernia Absence of contraindications for ketamine and opioids use patients with American Society of Anesthesiologist (ASA) physical status I,II

Exclusion criteria:

History of allergic reactions to the drug under study History of underlying diseases (diabetes, peptic ulcer, gastritis, coronary artery disease, renal failure) Any known psychiatric disorder Patients with incarcerated hernia patients with severe infection

Age

From **30 years** old to **60 years** old

Gender

Both

Phase

3

Groups that have been masked

- Participant
- Care provider

Sample size

Target sample size: **60**

Randomization (investigator's opinion)

Randomized

Randomization description

Patients are randomly assigned to two ketamine and fentanyl (control) groups using Random Allocation computer software. In this way, numbers are given to the patients according to the determined sample volume, and then the software allocates the patients to two ketamine and fentanyl (control) groups based on the given numbers.

Blinding (investigator's opinion)

Double blinded

Blinding description

After introducing the study to the patients and explaining the benefits and harms of the possible intervention, written consent is obtained from the patients. Then the patients are randomly assigned to one of the intervention or control groups. Patients and nurses who take care of the patients during the trial are blinded to the intervention and control groups, so that none of the nurses and patients know which drug, ketamine or fentanyl, the patients have been under general anesthesia with.

Placebo

Not used

Assignment

Parallel

Other design features

Secondary Ids

empty

Ethics committees

1

Ethics committee

Name of ethics committee

Ethics committee of urmia University of Medical Sciences

Street address

West Azarbaijan University of Medical Sciences, Resalat Blvd, urmia

City

urmia

Province

West Azarbaijan

Postal code

5714783734

Approval date

2023-04-19, 1402/01/30

Ethics committee reference number

IR.UMSU.REC.1402.040

Health conditions studied

1

Description of health condition studied

inguinal hernia

ICD-10 code

K40.9

ICD-10 code description

Unilateral or unspecified inguinal hernia, without obstruction or gangrene

Primary outcomes

1

Description

intensity of pain

Timepoint

At 60, 90 minutes and 24 hours after surgery

Method of measurement

Based on numeric rating scale (NRS)

2

Description

The consumption of opioids for pain after surgery

Timepoint

Up to 24 hours after the surgery

Method of measurement

Based on milligrams

Secondary outcomes

empty

Intervention groups

1

Description

Intervention group: In ketamine group, Induction of anesthesia is done with 1 to 4 mg/kg of intravenous ketamine, and then in maintenance phase, intravenous perfusion of ketamine at a rate of 2 to 10 µg/kg/min will be prescribed.

Category

Treatment - Drugs

2

Description

Control group: In fentanyl group, induction of anesthesia is performed with 1-2 µg/kg fentanyl as a routine anesthetic drug, and then maintenance phase, continuous infusion of fentanyl at a rate of 1-2 µg/kg/hr will be prescribed.

Category

Treatment - Drugs

Recruitment centers

1

Recruitment center

Name of recruitment center

Urmia Imam Khomeini educational and medical center

Full name of responsible person

Ali Akbar Nasiri

Street address

Imam Khomeini Educational and Medical Center, Ershad St., Urmia

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Sponsors / Funding sources

1

Sponsor

Name of organization / entity

Oroumia University of Medical Sciences

Full name of responsible person

Dr. Saber Gholizadeh

Street address

Research and Technology Vice-Chancellor, West Azarbaijan University of Medical Sciences and Healthcare Services, Emergency Room, Resalat Blvd., Urmia

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Grant name

Urmia University of Medical Sciences

Grant code / Reference number

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

Yes

Title of funding source

Oroumia University of Medical Sciences

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

Academic

Person responsible for general inquiries

Contact

Name of organization / entity

Oroumia University of Medical Sciences

Full name of responsible person

Ali Akbar Nasir

Position

Associate professor

Latest degree

Subspecialist

Other areas of specialty/work

Anesthesiology

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Person responsible for updating data

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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