

Clinical Trial Protocol

Iranian Registry of Clinical Trials

29 Jun 2026

Comparison intravenous anesthesia with Propofol and inhalation anesthesia with Isoflurane for pain measurement in post operative patients after inguinal herniorrhaphy

Protocol summary

Summary

The aim of this study is comparison intravenous anesthesia with Propofol and inhalation anesthesia with Isoflurane for pain measurement in post operative patients after inguinal herniorrhaphy. This study have a clinical trial design. Researcher chosen 100 patients in elective surgery of inguinal hernia. Vital sign such as heart rate, systolic and diastolic blood pressure and oxygen saturation measured before intubation of anesthesia. Researcher used 0/5 mg / kg Midazolam and 1 µg / kg Fentanyl for all patients. Researcher injected 2 mg / kg Propofol and 0/5 kg intravenous Atracurium in both groups. Patients randomly were divided in to two groups. The first group injected 100 mg /kg/minute Propofol for the maintenance of anesthesia and second group injected 1 mg/kg/minute Isoflurane for the maintenance of anesthesia. Intubation time and recovery time is recorded. Pain was measured in the recovery room (zero time) and after 6 hours (every 2 hours) based on the VAS scale. Patient's vital signs measured at zero time. Nausea and vomiting and shivering of patients were recorded during this period. 75 mg Diclofenac was injected for patient in the postoperative room if VAS scale was more than 3 during 6 hours after surgery .Dose of Diclofenac was recorded.

General information

Acronym

IRCT registration information

IRCT registration number: **IRCT2015112925277N1**
Registration date: **2015-12-24, 1394/10/03**
Registration timing: **registered_while_recruiting**

Last update:

Update count: **0**

Registration date

2015-12-24, 1394/10/03

Registrant information

Name

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Name of organization / entity

Arak University of Medical Sciences and Health Services

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Recruitment status

Recruitment complete

Funding source

Arak University of Medical Sciences and Health Services

Expected recruitment start date

2015-11-30, 1394/09/09

Expected recruitment end date

2017-02-27, 1395/12/09

Actual recruitment start date

empty

Actual recruitment end date

empty

Trial completion date

empty

Scientific title

Comparison intravenous anesthesia with Propofol and inhalation anesthesia with Isoflurane for pain measurement in post operative patients after inguinal herniorrhaphy

Public title

Comparison intravenous anesthesia with Propofol and inhalation anesthesia with Isoflurane for pain

measurement in post operative patients after inguinal herniorrhaphy

Purpose

Treatment

Inclusion/Exclusion criteria

Exclusion criteria: history of allergic reaction to the drug used in this study, pregnancy, drug addiction, and pain relief medications 24 hours before surgery, persistent hypertension, cardiovascular disease, renal failure

Age

No age limit

Gender

Both

Phase

N/A

Groups that have been masked

No information

Sample size

Target sample size: 100

Randomization (investigator's opinion)

Randomized

Randomization description**Blinding (investigator's opinion)**

Double 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 Arak University of Medical Sciences and Health Services

Street address

Research Center, Educational complex Payambar Azam, Basij square, Sardasht

City

Arak

Postal code

38481-7-6941

Approval date

2015-07-27, 1394/05/05

Ethics committee reference number

IR.ARAKMU.REC.1394.110

Health conditions studied**1****Description of health condition studied**

Inguinal herniorrhaphy

ICD-10 code

K40.3

ICD-10 code description

Unilateral or unspecified inguinal hernia, with obstruction, without gangrene

Primary outcomes**1****Description**

Pain

Timepoint

recovery room (zero time) and after 6 hours (every 2 hours)

Method of measurement

Visual Analog Scale

2**Description**

Temperature

Timepoint

Before and after surgery

Method of measurement

Thermometer

3**Description**

Blood pressure

Timepoint

Before and after surgery

Method of measurement

Barometer

4**Description**

Heart rate

Timepoint

Before and after surgery

Method of measurement

Rate in minute

5**Description**

Respiratory

Timepoint

Before and after surgery

Method of measurement

Rate in minute

6**Description**

Intubation time

Timepoint

After surgery

Method of measurement

Minute

7

Description

Recovery time

Timepoint

After surgery

Method of measurement

Minute

Secondary outcomes

1

Description

Diclofenac

Timepoint

After surgery

Method of measurement

Drug dose

Intervention groups

1

Description

first group injected 100 mg /kg/minute propofol for the maintenance of anesthesia

Category

Treatment - Drugs

2

Description

second group injected 1 mg/kg/minute isoflurane for the maintenance of anesthesia

Category

Treatment - Drugs

Recruitment centers

1

Recruitment center

Name of recruitment center

Valiasr Hospital

Full name of responsible person

Tahereh Mohaqeq

Street address

Valiasr Hospital ,Arak

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Sponsors / Funding sources

1

Sponsor

Name of organization / entity

Vice Chancellor of Arak University of Medical Sciences and Health Services

Full name of responsible person

Dr Bijan Yazdi

Street address

Research Center, Educational complex Payambar Azam, Basij square, Sardasht

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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

Vice Chancellor of Arak University of Medical Sciences and Health Services

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

Contact

Name of organization / entity

Arak University of Medical Sciences and Health Services

Full name of responsible person

Tahereh Mohaqeq

Position

Resident of Anesthesia

Other areas of specialty/work

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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