

Clinical Trial Protocol

Iranian Registry of Clinical Trials

02 Jul 2026

The effect of preventive lidocaine treatment on postoperative pain control

Protocol summary

Summary

Nowadays, gynecologic surgeries are one of the most common procedures in the outpatient settings and are performed progressively with laparoscopy. However, acute pain is a most important postoperative complication and not only, can lead to patient discomfort and unsatisfying, but also result to delayed discharge, hospital admission and increased costs. Severity of post laparoscopic pain is moderate to severe and about 35% to 65% of patients experienced it as abdominal or shoulder tip pain and up to 80% of patients required analgesia. Lidocaine has antihyperalgetic and anti inflammatory effects and use of this drug in the perioperative period can be a useful approach for post laparoscopic pain control. After given informed consent, all patient's will randomly allocate in one of the lidocaine or placebo groups. With induction of anesthesia, lidocaine group will receive lidocaine infusion in the rate of 2mg/kg/h and placebo group will only receive normal saline in the same infusion rate. After completion of surgery and extubation, pain severity will be evaluate with visual analogue scale (VAS) scoring system in recovery and 2,4,6,12 and 24 hours postoperatively. Postoperative pain with VAS \geq 4 will be controlled with 0.5/kg intravenous Meperidine. Pain severity, time to first analgesic request, total analgesic consumption doses and any probable postoperative complications will be recorded and statistically analyzed.

General information

Acronym

IRCT registration information

IRCT registration number: **IRCT2014040511700N5**

Registration date: **2014-08-20, 1393/05/29**

Registration timing: **registered_while_recruiting**

Last update:

Update count: **0**

Registration date

2014-08-20, 1393/05/29

Registrant information

Name

Farnaz Moslemi Tabrizi

Name of organization / entity

Alzahra hospital, Tabriz university of Medical Sciences

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Recruitment status

Recruitment complete

Funding source

Vice Chancellor for research of Tabriz university of Medical Sciences

Expected recruitment start date

2014-04-05, 1393/01/16

Expected recruitment end date

2015-04-05, 1394/01/16

Actual recruitment start date

empty

Actual recruitment end date

empty

Trial completion date

empty

Scientific title

The effect of preventive lidocaine treatment on postoperative pain control

Public title

Evaluation of the effect of perioperative intravenous infusion of lidocaine in postoperative pain management in women under gynecologic laparoscopy

Purpose

Prevention

Inclusion/Exclusion criteria

Inclusion criteria: Women candidate for Gynecologic Laparoscopic surgery; patients with ASA physical status I and II Exclusion criteria: Patient's with ASA class III or above; women with cardiovascular; respiratory diseases; hepatic or renal dysfunction; endocrine disorders; hypertension; psychologic disorder; convulsion; allergy to lidocaine; patients who received opioid or non opioid analgesic preoperatively

Age

From **18 years** old to **60 years** old

Gender

Female

Phase

N/A

Groups that have been masked

No information

Sample size

Target sample size: **60**

Randomization (investigator's opinion)

Randomized

Randomization description

Blinding (investigator's opinion)

Double blinded

Blinding description

Placebo

Used

Assignment

Parallel

Other design features

Secondary Ids

empty

Ethics committees

1

Ethics committee

Name of ethics committee

Ethics Committee of Vice chancellor for research,
Tabriz University of Medical Sciences

Street address

Tabriz university of Medical Sciences, Golgasht Ave,
Tabriz

City

Tabriz

Postal code

Approval date

2014-02-12, 1392/11/23

Ethics committee reference number

92192

Health conditions studied

1

Description of health condition studied

Postoperative pain after laparoscopy

ICD-10 code

R50,R51,R5

ICD-10 code description

Acute pain

Primary outcomes

1

Description

Pain

Timepoint

2, 4, 6, 12, 24 hours postoperative

Method of measurement

Visual Analog Scale (VAS)

Secondary outcomes

1

Description

Rescue analgesic consumption after surgery

Timepoint

24 hours postoperative

Method of measurement

Total dose mg

2

Description

The first analgesia administration

Timepoint

The time to the first analgesia administration

Method of measurement

Hours

Intervention groups

1

Description

With induction of anesthesia, lidocaine group will receive lidocaine infusion in the rate of 2mg/kg/h

Category

Prevention

2

Description

Control group will only receive normal saline in the same volume and infusion rate.

Category

Treatment - Drugs

Recruitment centers

1

Recruitment center

Name of recruitment center

Alzahra Hospital

Full name of responsible person

Dr. Farnaz Moslemi

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Web page address**Person responsible for scientific inquiries****Contact****Name of organization / entity**

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Medical Sciences**Full name of responsible person**

dr.Hassan Solaimanpour

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Grant name**Grant code / Reference number****Is the source of funding the same sponsor organization/entity?**

Yes

Title of funding sourceVice Chancellor for research of Tabriz 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****Contact****Name of organization / entity**

Tabriz University of Medical Sciences

Full name of responsible person

Dr. farnaz Moslemi

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Associate Professors of Anesthesiology

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Web page address**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