

Clinical Trial Protocol

Iranian Registry of Clinical Trials

27 Jun 2026

The effect of general and epidural anesthesia on respiratory parameters during and after laparoscopic cholecystectomy surgery

Protocol summary

Summary

The present study aims to compare the effect of general and epidural anesthesia on respiratory parameters during and after laparoscopic cholecystectomy surgery. Samples include all patients receiving laparoscopic cholecystectomy surgery in Imam Reza University. It is a double-blind clinical trial. 58 patients are selected among all patients receiving laparoscopic cholecystectomy surgery, and are divided to two groups randomly, a group receiving general and the other receiving epidural anesthesia. After pre-medication among GA with midazolam 0.5 mg/kg, intravenous fentanyl 1 µg/kg, patients are anesthetized using propofol 2 mg/kg and atracurium 5.0 mg/kg and standard monitoring is completed. Proper endotracheal tube is contrived and surgery starts. During the surgery, anesthetization continues through using isoflurane 1.4 1% blended with a mixture of air and 50% oxygen. Epidural set is used to inject 10 cc Lidocaine and epinephrine 1 in 200000, including 1 cc sodium bicarbonate in the space between vertebrae of the thoracic among EA group. ABG is tested during and after surgery and VC and SPO2 is measure through monitoring. Two groups are compared later. Entrance criteria includes written consent by patients; I, II ASA class; age range of 20 to 60; no history on respiratory or cardio problems; no history on diabetes or similar diseases needing treatment. Exclusion criterion is a surgery lasting more than 45 minutes.

General information

Acronym

IRCT registration information

IRCT registration number: **IRCT2014081714333N19**

Registration date: **2014-09-07, 1393/06/16**

Registration timing: **registered_while_recruiting**

Last update:

Update count: **0**

Registration date

2014-09-07, 1393/06/16

Registrant information

Name

Feizollah Foroughi

Name of organization / entity

kermanshah University of Medical Sciences

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Iran (Islamic Republic of)

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Recruitment status

Recruitment complete

Funding source

Kermanshah University of Medical Sciences

Expected recruitment start date

2014-06-15, 1393/03/25

Expected recruitment end date

2014-09-16, 1393/06/25

Actual recruitment start date

empty

Actual recruitment end date

empty

Trial completion date

empty

Scientific title

The effect of general and epidural anesthesia on respiratory parameters during and after laparoscopic cholecystectomy surgery

Public title

The effect of general and epidural anesthesia on respiratory parameters during and after laparoscopic cholecystectomy surgery

Purpose

Treatment

Inclusion/Exclusion criteria

Entrance criteria includes written consent by patients; I, II ASA class; age range of 20 to 60; no history on respiratory or cardio problems; no history on diabetes or similar diseases needing treatment. Exclusion criterion is a surgery lasting more than 45 minutes.

Age

From **20 years** old to **60 years** old

Gender

Both

Phase

N/A

Groups that have been masked

No information

Sample size

Target sample size: **58**

Randomization (investigator's opinion)

Randomized

Randomization description

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

Vice Chancellor for Research Affairs, Kermanshah University of Medical Sciences

Street address

Building No.2, Shahid Beheshti, Vice Chancellor for Research Affairs, Kermanshah University of Medical Sciences

City

Kermanshah

Postal code

Approval date

2014-06-09, 1393/03/19

Ethics committee reference number

10218/420/7/پ

Health conditions studied

1

Description of health condition studied

Laparoscopic cholecystectomy

ICD-10 code

K80.2

ICD-10 code description

Cholecystolithiasis

Primary outcomes

1

Description

FEV1

Timepoint

A day before and after surgery

Method of measurement

Using spirometry

2

Description

VC

Timepoint

During and an hour after surgery

Method of measurement

Using anesthesia machine monitor

3

Description

SPO2

Timepoint

Before, during and an hour after surgery

Method of measurement

Using pulse oxy meter monitor

4

Description

PH

Timepoint

Before, during and an hour after surgery

Method of measurement

Using blood gas analysis (ABG)

5

Description

Pao2

Timepoint

Before, during and an hour after surgery

Method of measurement

Using blood gas analysis (ABG)

6

Description

PaCo2

Timepoint

Before, during and an hour after surgery

Method of measurement

Using blood gas analysis (ABG)

7

Description

Hco3

Timepoint

Before, during and an hour after surgery

Method of measurement

Using blood gas analysis (ABG)

8**Description**

FVC

Timepoint

A day before and after surgery

Method of measurement

Through spirometry

9**Description**

O2SAT

Timepoint

Before, during and an hour after surgery

Method of measurement

Using blood gas analysis (ABG)

Secondary outcomes

empty

Intervention groups**1****Description**

In the first group(GA), After pre-medication among GA with midazolam 05.0 mg/kg, intravenous fentanyl 1 µg/kg, patients are anesthetized using propofol 2 mg/kg and atracurium 5.0 mg/kg and standard monitoring is completed. Proper indotracheal tube is contrived and surgery starts. During the surgery, anesthetization continues through using isoflurane 1.4 1% blended with mixture of air and 50% oxygen. During the surgery ABG is tested and SPO2 is measured accordingly. An hour later, during recovery period, ABG is retested and VC and SPO2 are measured through monitoring.

Category

Treatment - Surgery

2**Description**

In the second group (EA), 10 cc Lidocaine and epinephrine 1 in 200000, including 1 cc sodium bicarbonate for each 10 cc of Lidocaine, would be injected in the space between vertebrae of the thoracic using epidural set. ABG and SPO2 would be measured during surgery. ABG would be measured again, and VC and SPO2 would be checked using monitor an hour after surgery during recovery period.

Category

Treatment - Surgery

Recruitment centers**1****Recruitment center****Name of recruitment center**

Emam Reza Hospital

Full name of responsible person

Dr.iraj Peiman

Street address

Emam Reza Hospital,Razi Boulevard,Sorkhleje

City

Kermanshah

Sponsors / Funding sources**1****Sponsor****Name of organization / entity**

Vice Chancellor for Research Affairs, Kermanshah University of Medical Sciences

Full name of responsible person

Korosh Hamzehee

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Building No.2, Shahid Beheshti, Vice Chancellor for Research Affairs, Kermanshah University of Medical Sciences

City

Kermanshah

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

Yes

Title of funding source

Vice Chancellor for Research Affairs, Kermanshah 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****Person responsible for scientific inquiries****Contact****Name of organization / entity**

Kermanshah University of Medical Sciences

Full name of responsible person

Dr. Ali karbasforushan

Position

Anesthesiologist

Other areas of specialty/work**Street address**

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Web page address**Person responsible for updating data****Contact****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