

# Clinical Trial Protocol

## Iranian Registry of Clinical Trials

10 Jun 2026

### The comparison of Propofol and Isoflurane on arterial oxygenation pressure, mean arterial pressure and heart rate variations following one-lung ventilation in thoracic surgeries

#### Protocol summary

##### Summary

Objectives: Determination and comparison of effect of Propofol and Isoflurane on arterial oxygenation pressure, mean arterial pressure and heart rate variations following one-lung ventilation in thoracic surgeries  
Design: Randomized, not-blinded, Target sample size 61 people. The major inclusion criteria: The age range of 18-75 years old; ASA class I (Healthy person) and III (Severe systemic disease); The consent to undergo thoracic surgery with OLV, according to the study conditions The major exclusion criteria: Liver malfunction (Aspartate transaminase (AST) more than 40, Alanine transaminase (ALT) more than 40); Ischemic or valvular heart disease (history of cardiovascular diseases, physical status, Electrocardiogram (EKG) and echocardiography of the patients were analyzed); Recent use of anesthetic medications; Obstructive and end-stage restrictive pulmonary disease; Patients with one-lung ventilation (OLV) less than 30 min; Patients with End-Tidal CO<sub>2</sub> (ETCO<sub>2</sub>) more than 45 mmHg, and the respiratory rate of 12 breaths per minute; Patients who have pathological problems in the left lung (pulmonary), according to High Resolution Computed Tomography (HRCT) and pulmonary function tests (PFT) before the surgery. Intervention group: Intervention group: propofol (100 µg/kg/min), During Surgery. Control group: Control group: one Mac (1.1%) isoflurane, During surgery. Primary outcome measure: mean pressure of arterial oxygen and Mean Arterial pressure and Heart rate, During two lung ventilation and in 5, 10 and 20 minute after one lung ventilation.

#### General information

##### Acronym

##### IRCT registration information

IRCT registration number: **IRCT2013122615942N1**

Registration date: **2014-02-03, 1392/11/14**

Registration timing: **retrospective**

Last update:

Update count: **0**

##### Registration date

2014-02-03, 1392/11/14

##### Registrant information

###### Name

Mohammad Reza Rhnama Zadeh

###### Name of organization / entity

Ghaem Hospital

###### Country

Iran (Islamic Republic of)

###### Phone

+98 51 1768 7898

###### Email address

rahnamazadehm891@mums.ac.ir

##### Recruitment status

**Recruitment complete**

##### Funding source

Vice Chancellor for Research of Mashhad University of Medical Sciences

##### Expected recruitment start date

2012-02-01, 1390/11/12

##### Expected recruitment end date

2013-02-01, 1391/11/13

##### Actual recruitment start date

empty

##### Actual recruitment end date

empty

##### Trial completion date

empty

##### Scientific title

The comparison of Propofol and Isoflurane on arterial

oxygenation pressure, mean arterial pressure and heart rate variations following one-lung ventilation in thoracic surgeries

#### Public title

Propofol and Isoflurane effects in thoracic surgeries

#### Purpose

Treatment

#### Inclusion/Exclusion criteria

The major inclusion criteria: The age range of 18-75 years old; ASA class I (Healthy person) and III (Severe systemic disease); The consent to undergo thoracic surgery with OLV, according to the study conditions The major exclusion criteria: Liver malfunction (Aspartate transaminase (AST) more than 40, Alanine transaminase (ALT) more than 40); Ischemic or valvular heart disease (history of cardiovascular diseases, physical status, Electrocardiogram (EKG) and echocardiography of the patients were analyzed); Recent use of anesthetic medications; Obstructive and end-stage restrictive pulmonary disease; Patients with one-lung ventilationin (OLV) less than 30 min; Patients with End -Tidal CO2 (ETCO2) more than 45 mmHg, and the respiratory rate of 12 breaths per minute; Patients who have pathological problems in the left lung (pulmonary), according to High Resolution Computed Tomography (HRCT) and pulmonary function tests (PFT) before the surgery.

#### Age

From **18 years** old to **75 years** old

#### Gender

Both

#### Phase

2

#### Groups that have been masked

*No information*

#### Sample size

Target sample size: **60**

#### 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

Ethics Committee of Mashhad University of Medical Sciences

###### Street address

Vice Chancellor for Research of Mashhad University of Medical Sciences, Ghoreishi Building, Daneshgah street, Mashhad

#### City

Mashhad

#### Postal code

#### Approval date

2013-02-02, 1391/11/14

#### Ethics committee reference number

910182

### Health conditions studied

#### 1

##### Description of health condition studied

Anasthesia

##### ICD-10 code

Y48.4

##### ICD-10 code description

Anaesthetic, unspecified

### Primary outcomes

#### 1

##### Description

mean pressure of arterial oxygen

##### Timepoint

During two lung ventilation and in 5, 10 and 20 minute after one lung ventilation

##### Method of measurement

ABG Test by GEM premier 3000

#### 2

##### Description

Mean Arterial pressure

##### Timepoint

During two lung ventilation and in 5, 10 and 20 minute after one lung ventilation

##### Method of measurement

ABG Test by GEM premier 3000

#### 3

##### Description

Heart Rate

##### Timepoint

During two lung ventilation and in 5, 10 and 20 minute after one lung ventilation

##### Method of measurement

ABG Test by GEM premier 3000

### Secondary outcomes

#### 1

##### Description

Arterial oxygen saturation

##### Timepoint

During ventilation of two lungs, 5, 10 and 20 minute after

one lung ventilation  
**Method of measurement**  
Monitoring

## Intervention groups

### 1

**Description**  
Intervention group: Propofol (100 µg/kg/min), During Surgery  
**Category**  
Prevention

### 2

**Description**  
Control group: one Mac (1.1%) Isoflurane, During surgery  
**Category**  
Prevention

## Recruitment centers

### 1

**Recruitment center**  
**Name of recruitment center**  
Ghaem Hospital  
**Full name of responsible person**  
Mohammad Reza Rahnamazadeh  
**Street address**  
Ghaem Hospital, Ahmadabad Street, Mashhad  
**City**  
Mashhad

## Sponsors / Funding sources

### 1

**Sponsor**  
**Name of organization / entity**  
Vice Chancellor for Research of Mashhad University of Medical Sciences  
**Full name of responsible person**  
Mohammad Ramezani  
**Street address**  
Vice Chancellor for Research of Mashhad University of Medical Sciences, Ghoreishi Building, Daneshgah Street, Mashhad  
**City**  
Mashhad  
**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 of Mashhad 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**  
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Ghaem Hospital  
**Full name of responsible person**  
Mohammad Reza Rahnamazadeh  
**Position**  
Student  
**Other areas of specialty/work**  
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## Person responsible for scientific inquiries

**Contact**  
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## Person responsible for updating data

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student

**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*