

Clinical Trial Protocol

Iranian Registry of Clinical Trials

03 Jul 2026

A Comparison and Evaluation of intra ocular pressure following intubation with Mccoy, Macintosh and video laryngoscope (Glide scope) in patients with cataract surgery.

Protocol summary

Summary

Our general objective in this study is assessing intraocular pressure (IOP) changes after intubation with three types of laryngoscope (McCoy, Macintosh and video laryngoscope or Glidescope). This is a double blind and prospective clinical trial on 180 patients subjected to cataract surgery in Shafa Hospital. Inclusion criteria are: patients who are admitted for elective cataract surgery, age: 20-70 year and have ASA classification as 1 or 2. Exclusion criteria are: history of alcohol and drug abuse, antihypertensive and cardiovascular medicines usage, glaucoma eye drop usage, diabetes and open eye injuries. The patients will be divided into three groups. Induction of anesthesia in all patients is similar. Just after induction and before intubation, intraocular pressure (IOP) will be measured by means of Schiötz tonometer. Then intubation will be done by McCoy laryngoscope in group A, Macintosh laryngoscope in group B and glidescope in group C. After intubation IOP is measured again and anesthesia is maintained on Isoflurane. IOP is measured again five minutes later. Collected data will be analyzed by Repeated Measures ANOVA test.

General information

Acronym

IRCT registration information

IRCT registration number: **IRCT201301079585N3**
Registration date: **2013-02-21, 1391/12/03**
Registration timing: **retrospective**

Last update:

Update count: **0**

Registration date

2013-02-21, 1391/12/03

Registrant information

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Name of organization / entity

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Recruitment status

Recruitment complete

Funding source

Vice chancellor for research, Kerman University of Medical Sciences

Expected recruitment start date

2012-08-22, 1391/06/01

Expected recruitment end date

2013-01-20, 1391/11/01

Actual recruitment start date

empty

Actual recruitment end date

empty

Trial completion date

empty

Scientific title

A Comparison and Evaluation of intra ocular pressure following intubation with McCoy, Macintosh and video laryngoscope (Glide scope) in patients with cataract surgery.

Public title

Evaluation of intra ocular pressure following intubation

Purpose

Treatment

Inclusion/Exclusion criteria

Inclusion criteria are: patients who are admitted for

elective cataract surgery, age: 20-70 and have ASA classification as 1 or 2. Exclusion criteria are: history of alcohol and drug abuse, antihypertensive and cardiovascular medicines usage, glaucoma eye drop usage, diabetes and open eye injuries.

Age

From **20 years** old to **70 years** old

Gender

Both

Phase

N/A

Groups that have been masked

No information

Sample size

Target sample size: **180**

Randomization (investigator's opinion)

Randomized

Randomization description

Blinding (investigator's opinion)

Double blinded

Blinding description

Placebo

Not used

Assignment

Parallel

Other design features

All patients will sign informed consent and they will be divided into three groups by means of random numbers table. Intubation is done by an anesthesiologist and the resident of anesthesiology who is blind to the kind of laryngoscope will obtain IOP, heart rate and blood pressure. Moreover the patients are unaware of the kind of used laryngoscope.

Secondary Ids

empty

Ethics committees

1

Ethics committee

Name of ethics committee

Ethnic Committee of Kerman University of Medical Sciences

Street address

Jomhoori Boulevard

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Postal code

Approval date

2012-04-17, 1391/01/29

Ethics committee reference number

k/91/02

Health conditions studied

1

Description of health condition studied

cataract

ICD-10 code

H26.9

ICD-10 code description

Cataract, unspecified

Primary outcomes

1

Description

intra ocular pressure

Timepoint

before and after intubation, and 5 minutes later

Method of measurement

Schiotz tonometer

Secondary outcomes

1

Description

Mean Arterial Pressure

Timepoint

Before and after intervention and every 5 minutes during operation

Method of measurement

mmhg / by electronic monitoring

2

Description

Heart Rate

Timepoint

Before and after intervention and every 5 minutes during operation

Method of measurement

electronic monitoring

Intervention groups

1

Description

Intubation by Mccooy laryngoscope made by Riester company of Germany, after induction of anesthesia.

Category

Treatment - Devices

2

Description

Intubation by Macintosh laryngoscope made by Riester company of Germany, after induction of anesthesia.

Category

Treatment - Devices

3

Description

Intubation by glidescope made by Verathon® Medical (Canada) ULC, after induction of anesthesia.

Category

Treatment - Devices

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

Shafa Hospital

Full name of responsible person

Zahra Asskari

Street address

Shafa Hospital, Shafa Street

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

Vice Chancellor for research, Kerman University of Medical Sciences

Full name of responsible person

Fatemeh Hassani

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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 for research, Kerman 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**

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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