

Clinical Trial Protocol

Iranian Registry of Clinical Trials

17 Sep 2026

Comparing isoflurane with sevoflurane anesthesia maintenance on recovery of out patient pediatric surgery with laryngeal mask

Protocol summary

Study aim

Comparing of Isoflurane with Sevoflurane anesthesia on recovery in children

Design

Clinical trial with intervention groups, sample size of 40 patients per group, with parallel groups, double-blinded, randomized

Settings and conduct

In this double-blind randomized clinical trial, 80 patients from Tabriz children hospital with American society of Anesthesiologists (ASA) status I and II, undergoing elective outpatient operation under an hour (from March 2019 to August 2019) will be enrolled. The patients will be randomized in one of the two groups of Isoflurane and sevoflurane anesthesia.

Participants/Inclusion and exclusion criteria

Inclusion criteria: children between 2 months- 6 years old Indicative Outpatient Surgeries admitted to the Children's Hospital Parental informed consent for the study ASA class I and II Exclusion criteria: Sensitivity to any of the drugs used in the study Children with underlying illnesses such as diabetes; asthma; congenital heart disease and ...

Intervention groups

Intervention group1: Inhalation anesthesia with sevoflurane. received midazolam (0.03-0.05 mg/kg), Fentanyl 1µg/kg as a premedication. Anesthesia induction will be with MAC %4-5 Sevoflurane , 50% N2O, 50% O2 .LMA will be inserted with the right size. Anesthesia maintenance will be with Sevoflurane inhalation 2-3% , 50% N2O, 50% O2 Intervention group2: Inhalation anesthesia with sevoflurane. received midazolam (0.03-0.05 mg/kg), Fentanyl 1µg/kg as a premedication. . Anesthesia induction will be with MAC %2.5 - 3 Isoflurane , 50% N2O, 50% O2 .LMA will be inserted with the right size. Anesthesia maintenance will be with Isoflurane inhalation 1.2%-1.5% , 50% N2O, 50% O2

Main outcome variables

duration of stay in recovery, Blood pressure, Heart rate, pain, nausea

General information

Reason for update

Acronym

IRCT registration information

IRCT registration number: **IRCT20100527004041N19**

Registration date: **2019-07-30, 1398/05/08**

Registration timing: **registered_while_recruiting**

Last update: **2019-07-30, 1398/05/08**

Update count: **0**

Registration date

2019-07-30, 1398/05/08

Registrant information

Name

Mahin Seyedhejazi

Name of organization / entity

Tabriz University of Medical Sciences

Country

Iran (Islamic Republic of)

Phone

+98 41 3385 2252

Email address

seidhejazie@tbzmed.ac.ir

Recruitment status

Recruitment complete

Funding source

Expected recruitment start date

2019-07-23, 1398/05/01

Expected recruitment end date

2020-08-20, 1399/05/30

Actual recruitment start date

empty

Actual recruitment end date

empty

Trial completion date
empty

Scientific title
Comparing isoflurane with sevoflurane anesthesia maintenance on recovery of out patient pediatric surgery with laryngeal mask

Public title
Comparing of recovery isoflurane with sevoflurane anesthesia in children

Purpose
Treatment

Inclusion/Exclusion criteria
Inclusion criteria:
 children between 2 months - 6 years old Indicative Outpatient Surgeries admitted to the Children's Hospital Parental informed consent for the study ASA class I and II
Exclusion criteria:
 Sensitivity to any of the drugs used in the study Children with underlying illnesses such as diabetes; asthma; congenital heart disease and ...

Age
From **2 months** old to **6 years** old

Gender
Both

Phase
2-3

Groups that have been masked

- Participant
- Investigator

Sample size
Target sample size: **80**

Randomization (investigator's opinion)
Randomized

Randomization description
Random method: Block, Random unit: individual, Random Tool: Random Block 4. for this purpose, 25 blocks with 4 subjects in each block will be used. the combination of all patterns will be considered including AABB, ABAB, BABA, BBAA, BABA, BAAB. For selecting each blocks, dice drooped and the block number will be selected. this procedure continued to completed the allocation and reached to sample size.

Blinding (investigator's opinion)
Double blinded

Blinding description
This is a double-blinded study and Researcher and patients are kept unaware of intervention in each group and In order to allocation concealment, the type of intervention will be written on paper and placed in numbered, matte and packed envelopes. The envelopes will be opened in the order of participation of the participants and the type of group will be determined.

Placebo
Not used

Assignment
Parallel

Other design features

Secondary Ids

empty

Ethics committees

1

Ethics committee

Name of ethics committee

Ethic Committee of Tabriz University of Medical Sciences

Street address

Tabriz University of Medical Science, Gholghasht street, Azadi street, Tabriz

City

Tabriz

Province

East Azarbaijan

Postal code

5166616471

Approval date

2019-05-09, 1398/02/19

Ethics committee reference number

IR.TBZMED.REC.1398.126

Health conditions studied

1

Description of health condition studied

General Anesthesia

ICD-10 code

Y48.2

ICD-10 code description

Other and unspecified general anaesthetics

Primary outcomes

1

Description

Duration of stay in recovery

Timepoint

After entering till release from recovery

Method of measurement

According to Alderet Score

2

Description

Blood pressure

Timepoint

Every 20 minutes till release from recovery

Method of measurement

Monitor

3

Description

Heart Rate

Timepoint

Every 20 minutes till release from recovery

Method of measurement

ECG device

4

Description

Nausea

Timepoint

After entering till release from recovery

Method of measurement

Visual Analogue Scale

5

Description

pain

Timepoint

After entering till release from recovery

Method of measurement

wong - baker pain scale

Secondary outcomes

empty

Intervention groups

1

Description

Intervention group1: received midazolam 0.03-0.05 mg/kg and Fentanyl 1µg/kg as a premedication. Anesthesia induction will be with MAC %4-5 Sevoflurane , 50% N2O, 50% O2 .LMA will be inserted with the right size. Anesthesia maintenance will be with Sevoflurane inhalation 2-3% , 50% N2O, 50% O2

Category

Treatment - Drugs

2

Description

Intervention group2: received midazolam 0.03-0.05 mg/kg and Fentanyl 1µg/kg as a premedication. Anesthesia induction will be with MAC %2.5 - 3 Isoflurane, 50% N2O, 50% O2. LMA will be inserted with the right size. Anesthesia maintenance will be with Isoflurane inhalation 1.2%-1.5% , 50% N2O, 50% O2

Category

Treatment - Drugs

Recruitment centers

1

Recruitment center

Name of recruitment center

Tabriz Children Hospital

Full name of responsible person

Dr.Mahin Seyedhejazi

Street address

Children's Hospital, Sheshgelan st

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seidhejazie@tbzmed.ac.ir

Web page address

<https://childrenhospital.tbzmed.ac.ir>

Sponsors / Funding sources

1

Sponsor

Name of organization / entity

Tabriz University of Medical Sciences

Full name of responsible person

Abolghasem Jouyban

Street address

Research and innovation deputy, third floor, No 2
Central Building, Tabriz University of Medical
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Web page address

<https://www.tbzmed.ac.ir>

Grant name

Grant code / Reference number

Is the source of funding the same sponsor organization/entity?

Yes

Title of funding source

Tabriz University of Medical Sciences

Proportion provided by this source

100

Public or private sector

Public

Domestic or foreign origin

Domestic

Category of foreign source of funding

empty

Country of origin

Type of organization providing the funding

Academic

Person responsible for general inquiries

Contact

Name of organization / entity

Tabriz University of Medical Sciences

Full name of responsible person

Afsaneh Parvize

Position

Assistant of Anesthesiology

Latest degree

Specialist

Other areas of specialty/work

Anesthesiology

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Name of organization / entity

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Full name of responsible person

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Position

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Latest degree

Specialist

Other areas of specialty/work

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Person responsible for scientific inquiries

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Tabriz University of Medical Sciences

Full name of responsible person

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Position

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Latest degree

Specialist

Other areas of specialty/work

Anesthesiology

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Sharing plan

Deidentified Individual Participant Data Set (IPD)

Yes - There is a plan to make this available

Study Protocol

Yes - There is a plan to make this available

Statistical Analysis Plan

Yes - There is a plan to make this available

Informed Consent Form

Yes - There is a plan to make this available

Clinical Study Report

Yes - There is a plan to make this available

Analytic Code

No - There is not a plan to make this available

Data Dictionary

Undecided - It is not yet known if there will be a plan to make this available

Title and more details about the data/document

A portion of the data that represents the final outcome

When the data will become available and for how long

Access will be 6 months after the results are printed.

To whom data/document is available

All Physicians and residents of the department of Anesthesia

Under which criteria data/document could be used

After obtaining permission from the deputy research of group

From where data/document is obtainable

Person responsible for scientific accountability of study

What processes are involved for a request to access data/document

First approved by the Research Vice-President

Comments

Person responsible for updating data

Contact