

# Clinical Trial Protocol

## Iranian Registry of Clinical Trials

10 Jun 2026

### Comparing of recovery in total intravenous anesthesia(TIVA) with inhalation anaesthetic (sevoflurane) in children between 2-10 years old

#### Protocol summary

##### Study aim

Comparing of recovery in total intravenous anesthesia(TIVA) with inhalation anaesthetic (sevoflurane) in children

##### Design

Clinical trial with intervention groups, 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 total intravenous(T) and sevoflurane anesthesia.

##### Participants/Inclusion and exclusion criteria

inclusion criteria: children between 2-10 years old, Indicative Outpatient Surgery, 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:" Total Intravenous Anesthesia.received midazolam (0/03-0/05 mg/kg), Fentanyl 1µg/kg as a premedication. Anesthesia induction will be with Lidocaine 1mg / kg and propofol 3-5 mg / kg , Then LMA will be inserted with the right size. . Anesthesia maintenance with total Intravenous Anesthesia will be by infusion of propofol 100-200 mcg / kg and remifentanyl 0.1 mcg / kg, 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 Sevoflurane 8% , 50% N2O, 50% O2 .LMA will be inserted with the right size. Anesthesia maintenance will be by Sevoflurane inhalation 2-3% , 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: **IRCT20190114042358N1**

Registration date: **2019-02-25, 1397/12/06**

Registration timing: **registered\_while\_recruiting**

Last update: **2019-02-25, 1397/12/06**

Update count: **0**

##### Registration date

2019-02-25, 1397/12/06

##### Registrant information

##### Name

Daryoush Sheikhzadeh

##### Name of organization / entity

##### Country

Iran (Islamic Republic of)

##### Phone

+98 41 3526 2257

##### Email address

dr.d.sheik@gmail.com

##### Recruitment status

**Recruitment complete**

##### Funding source

##### Expected recruitment start date

2019-02-20, 1397/12/01

##### Expected recruitment end date

2019-08-21, 1398/05/30

##### Actual recruitment start date

empty

##### Actual recruitment end date

empty

**Trial completion date**  
empty

**Scientific title**  
Comparing of recovery in total intravenous anesthesia(TIVA) with inhalation anaesthetic (sevoflurane) in children between 2-10 years old

**Public title**  
Comparing of recovery in total intravenous anesthesia(TIVA) with inhalation anaesthetic (sevoflurane) in children

**Purpose**  
Treatment

**Inclusion/Exclusion criteria**  
**Inclusion criteria:**  
children between 2-10 years old Indicative Outpatient Surgery 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 years** old to **10 years** old

**Gender**  
Both

**Phase**  
2-3

**Groups that have been masked**

- Participant
- Outcome assessor

**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

2019-01-13, 1397/10/23

#### Ethics committee reference number

IR.TBZMED.REC.1397.834

## Health conditions studied

### 1

#### Description of health condition studied

inhalation anesthesia

#### ICD-10 code

Y48.0

#### ICD-10 code description

Inhaled anaesthetics

### 2

#### Description of health condition studied

Intravenous Anesthesia

#### ICD-10 code

Y48.1

#### ICD-10 code description

Parenteral 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 5 minutes till release from recovery

**Method of measurement**

Monitor

**3**

**Description**

Heart rate

**Timepoint**

Every 5 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 group:" Total Intravenous Anesthesia.received midazolam (0/03-0/05 mg/kg), Fentanyl 1µg/kg as a premedication. Anesthesia induction will be with Lidocaine 1mg / kg and propofol 3-5 mg / kg , Then LMA will be inserted with the right size. Anesthesia maintenance with total Intravenous Anesthesia is done by infusion of propofol 100-200 mcg / kg and remifentanil 0.1 mcg / kg, 50% N2O, 50% O2

**Category**

Treatment - Drugs

**2**

**Description**

"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 Sevoflurane 8% , 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

**Recruitment centers**

**1**

**Recruitment center**

**Name of recruitment center**

Tabriz Children's Hospital

**Full name of responsible person**

Daryoush Sheikhzadeh

**Street address**

Tabriz Children's Hospital, Sheshgelan St

**City**

Tabriz

**Province**

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**Postal code**

5136735886

**Phone**

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**Email**

dr.d.sheik@gmail.com

**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**

Tabriz University of Medical Sciences, Golghasht St

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ajouyban@hotmail.com

**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**

Daryoush Sheikhzadeh

**Position**

Assistant Professor

**Latest degree**

Specialist

**Other areas of specialty/work**

Anesthesiology

**Street address**

Tabriz Children's Hospital, Sheshgelan St

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**Web page address**

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

Tabriz University of Medical Sciences

**Full name of responsible person**

Daryoush Sheikhzadeh

**Position**

Assistant Professor

**Latest degree**

Specialist

**Other areas of specialty/work**

Anesthesiology

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**Person responsible for updating data****Contact****Name of organization / entity**

Tabriz University of Medical Sciences

**Full name of responsible person**

Mehdi Razaghipour Sorkhab

**Position**

Resident of Anesthesiology

**Latest degree**

Medical doctor

**Other areas of specialty/work**

Anesthesiology

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**Email**

Razagh92tabriz@yahoo.com

**Sharing plan****Deidentified Individual Participant Data Set (IPD)**

Yes - There is a plan to make this available

**Study Protocol**

No - There is not 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**

Undecided - It is not yet known if there will be 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**