

Clinical Trial Protocol

Iranian Registry of Clinical Trials

05 Jun 2026

Comparison of the effects of sevoflurane with Remifentanil plus Propofol on recovery conditions in adolescent patient with movement disorders candidate for dental treatment under general anesthesia.

Protocol summary

Study aim

Determining comparative study of recovery conditions in adolescent patient with movement disorders under general anesthesia with two pharmacological diets of sevoflurane with remifentanil plus propofol

Design

Clinical trial with control group, with parallel groups, double-blind, randomized, on 30 patients. The random number table is used for randomization.

Settings and conduct

The person evaluating the recovery conditions and the participants are unaware of the type of drug injected. For this purpose, the injector and the person evaluating the recovery conditions are different. This study is done in the operating room of the Faculty of Dentistry of Isfahan University of Medical Sciences. Adult patients with movement disorders to perform dental work under general anesthesia, after clinical evaluation by an anesthesiologist and filling out a general information questionnaire and consent form, enter the study. Using a Random number table, they fall into the category of intervention or control. Recovery conditions are then assessed with the "Nursing Delirium Screening Scale" checklist.

Participants/Inclusion and exclusion criteria

Inclusion criteria: The desire to participate in the study, Patients over 18 years of age, Patients with movement disorders, Patients are candidates for dental treatment under general anesthesia Exclusion criteria: People who are allergic to eggs or soy, Known allergies, Chronic diseases including heart disease, bleeding disorders, liver disease and kidney disease

Intervention groups

Intervention group: To maintain anesthesia, sevoflurane drug with MAC: 1.85 is used. Control group: To maintain anesthesia, Rimifentanil is taken with Propofol (Propofol: 3 mg / kg body weight, followed by Remifentanil: 1 µg /

kg body weight per minute).

Main outcome variables

Recovery period; The amount of delirium

General information

Reason for update

Acronym

IRCT registration information

IRCT registration number: **IRCT20100621004224N38**

Registration date: **2020-05-13, 1399/02/24**

Registration timing: **prospective**

Last update: **2020-05-13, 1399/02/24**

Update count: **0**

Registration date

2020-05-13, 1399/02/24

Registrant information

Name

Nasser Kaviani

Name of organization / entity

Country

Iran (Islamic Republic of)

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+98 31 1792 2858

Email address

kaviani@dnt.mui.ac.ir

Recruitment status

Recruitment complete

Funding source

Expected recruitment start date

2020-05-26, 1399/03/06

Expected recruitment end date

2020-06-19, 1399/03/30

Actual recruitment start date

empty

Actual recruitment end date

empty

Trial completion date

empty

Scientific title

Comparison of the effects of sevoflurane with Remifentanil plus Propofol on recovery conditions in adolescent patient with movement disorders candidate for dental treatment under general anesthesia.

Public title

Effects of sevoflurane with Remifentanil plus Propofol on recovery conditions

Purpose

Supportive

Inclusion/Exclusion criteria**Inclusion criteria:**

The desire to participate in the study Patients over 18 years of age Patients with movement disorders Patients are candidates for dental treatment under general anesthesia

Exclusion criteria:

People who are allergic to eggs or soy Known allergies Chronic diseases including heart disease, bleeding disorders, liver disease and kidney disease

Age

From **18 years** old

Gender

Both

Phase

3

Groups that have been masked

- Participant
- Outcome assessor

Sample size

Target sample size: **30**

Randomization (investigator's opinion)

Randomized

Randomization description

Balanced block randomization

Blinding (investigator's opinion)

Double blinded

Blinding description

This study is double blind and the injecting person and the person evaluating the recovery conditions are different. For this purpose, an out-of-study person is used to assess recovery conditions. The person evaluating the recovery conditions is not aware of the medication used for anesthesia. Participants are also unaware of the type of drug injected

Placebo

Not used

Assignment

Parallel

Other design features**Secondary Ids**

empty

Ethics committees**1****Ethics committee****Name of ethics committee**

Ethics Committee of Isfahan University of Medical Sciences

Street address

Faculty of Dentistry, Isfahan University of Medical Sciences, Hezarjerib Ave., Isfahan Town,

City

Isfahsn

Province

Isfahan

Postal code

8174673461

Approval date

2020-02-23, 1398/12/04

Ethics committee reference number

IR.MUI.RESEARCH.REC.1398.756

Health conditions studied**1****Description of health condition studied**

Movement disorders

ICD-10 code**ICD-10 code description****Primary outcomes****1****Description**

Recovery score on "Nursing Delirium Screening Scale" Checklist

Timepoint

5,10 and 15 minuts after entering recovery

Method of measurement

"Nursing Delirium Screening Scale" Checklist

Secondary outcomes

empty

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

Intervention group: to maintain anesthesia, sevoflurane USP inhaled drug from Piramal Company is used. The drug is given to the patient at a dose of MAC: 1.85 through a tracheal tube with a vaporizer for the drug sevoflurane.

Category

Other

2

Description

Control group: To maintain anesthesia, Rimifentanil is taken with Propofol (Propofol: 3 mg / kg body weight, followed by Remifentanil: 1 µg / kg body weight per minute). Depending on whether the patient is shaking or shows signs of alertness in response to the situation, further injection of the drug (remifentanil: 0.5 µg per kg body weight or propofol 0.5-1 mg / kg body weight) is done. The drugs are injected within 10-15 seconds.

Category

Other

Recruitment centers

1

Recruitment center

Name of recruitment center

The operating room of the Faculty of Dentistry, Isfahan University of Medical Sciences

Full name of responsible person

Naser Kaviani

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Faculty of Dentistry, Isfahan University of Medical Sciences, Hezarjerib Ave., Isfahan Town,

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Sponsors / Funding sources

1

Sponsor

Name of organization / entity

Esfahan University of Medical Sciences

Full name of responsible person

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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

Esfahan 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

Esfahan University of Medical Sciences

Full name of responsible person

Naser Kaviani

Position

Consultant

Latest degree

Ph.D.

Other areas of specialty/work

Anesthesiology

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Person responsible for scientific inquiries

Contact

Name of organization / entity

Esfahan University of Medical Sciences

Full name of responsible person

Naser Kaviani

Position

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Latest degree

Ph.D.

Other areas of specialty/work

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Person responsible for updating data

Contact

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Sharing plan

Deidentified Individual Participant Data Set (IPD)

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

Study Protocol

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

Statistical Analysis Plan

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

Informed Consent Form

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

Clinical Study Report

Undecided - It is not yet known if there will be 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