

Clinical Trial Protocol

Iranian Registry of Clinical Trials

10 Jun 2026

Comparison of the effect of intravenous ketamine and midazolam as a premedication in children undergoing cochlear implantation at the time of separation from parents

Protocol summary

Study aim

Inducing sedation for deaf and muteness children when leaving their parents for cochlear implantation.

Design

This prospective clinical study with parallel groups, double-blinded, will be performed on 74 children who will be referred to Baqiyatallah Hospital for cochlear implantation. Patients will be divided randomly into two groups of 37 patients, midazolam and ketamine.

Settings and conduct

This double-blind study will be conducted on children who will be referred to Baqiyatallah hospital for cochlear implantation. Patients will be randomly divided into midazolam and ketamine groups. The researcher's assistant who will be unaware of the type of used drugs will observe the child in the pre-operative room and fill out the SASS rating form for anxiety scoring. Then, another person will inject the medicine that will be encoded before (blindness will be kept). After calming the child, the researcher's assistant will fill out the form again by removing the child from the parent and inducing anesthesia.

Participants/Inclusion and exclusion criteria

Children aged 6 months to 5 years old in Class 1 and 2 of the American Anesthesiology Association will be included in the study. Children with known sensitivity to ketamine or midazolam, history of using narcotics, age less than 6 months, increased intracranial pressure, history of seizure, upper and lower respiratory tract infection, will be excluded.

Intervention groups

The group of midazolam (receiving 0.1 milligram per kilogram of midazolam produced by Tehran Chemistry Factory diluted with distilled water) and the ketamine group (1 milligram per kilogram of ketamine produced by Rotex medica diluted with distilled water)

Main outcome variables

The child's sedation when separating from the parents, the child's sedation during induction of anesthesia and child sedation in recovery room.

General information

Reason for update

Acronym

IRCT registration information

IRCT registration number: **IRCT20161022030421N3**

Registration date: **2019-06-15, 1398/03/25**

Registration timing: **prospective**

Last update: **2019-06-15, 1398/03/25**

Update count: **0**

Registration date

2019-06-15, 1398/03/25

Registrant information

Name

Marzieh Lak

Name of organization / entity

Country

Iran (Islamic Republic of)

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+98 21 2244 9013

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Recruitment status

Not yet recruiting

Funding source

Expected recruitment start date

2640-01-21, 2018/11/01

Expected recruitment end date

2640-05-22, 2019/03/01

Actual recruitment start date

empty

Actual recruitment end date

empty

Trial completion date

empty

Scientific title

Comparison of the effect of intravenous ketamine and midazolam as a premedication in children undergoing cochlear implantation at the time of separation from parents

Public title

Comparison of the effect of intravenous ketamine and midazolam as a premedication in children .

Purpose

Treatment

Inclusion/Exclusion criteria**Inclusion criteria:**

Children from 6 months to 5 years of age with ASA class I and II, candidate for cochlear implant

Exclusion criteria:

Children with known sensitivity to ketamine or midazolam, as well as history of taking psychiatric medications, age less than 6 months, increased ICP, brain tumor masses, history of seizure, upper and lower respiratory tract infection.

Age

From **6 months** old to **5 years** old

Gender

Both

Phase

3

Groups that have been masked

- Participant
- Care provider
- Investigator
- Outcome assessor

Sample size

Target sample size: **74**

Randomization (investigator's opinion)

Randomized

Randomization description

Patients will be randomly assigned into two groups of midazolam and ketamine. The person who codes the drug, for example, may will allocate even numbers to ketamine and odd numbers to midazolam, or vice versa.

Blinding (investigator's opinion)

Double blinded

Blinding description

The medication will be prepared by a person in the volume of 5 cc and the drug code is written on it. Then another person who will not know the type of medicine will inject the drug into the child and fill out the questionnaire and record the drug code in the questionnaire. The child and parents will not be aware of the type of medicine.

Placebo

Not used

Assignment

Parallel

Other design features**Secondary Ids**

empty

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

Ethics committee of Baqiyatallah University of Medical Sciences

Street address

Baqiyatallah University of Medical Science, Mollasadra Ave, Tehran, Iran

City

Tehran

Province

Tehran

Postal code

1435915371

Approval date

2018-10-29, 1397/08/07

Ethics committee reference number

IR.BMSU.REC.1397.031

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

Seduction of the chorus and dumb children

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

sedation when separating from parents

Timepoint

In the pre-operative room, when the child is separated from the parents

Method of measurement

Observe and filling the form of sedation anxiety score scale (SASS)

2**Description**

Sedation of the child during induction of anesthesia

Timepoint

On the operating room bed, Induction of anesthesia

Method of measurement

Using the form of sedation anxiety score scale (SASS)

3**Description**

Child sedation in recovery

Timepoint

From the child's arrival to the recovery until the release of the recovery

Method of measurement

Using PACU sedation score scale (PACUSSS)

Secondary outcomes

empty

Intervention groups

1

Description

Intervention group (Midazolam): In this group, midazolam will be given with dose of one tenth of mg / kg of midazolamole in a 5-cc syringe and diluted into 5 cc distilled water. In pre-anesthetic room, the medicine will be injected intravenously, while the baby will be monitored for oxygenation.

Category

Rehabilitation

2

Description

Intervention group (Ketamin): In this group, ketamine with dose of 1 mg per kg will be injected into a 5-cc syringe and diluted into 5 cc distilled water. In pre-anesthetic room, the medicine will be injected intravenously, while the baby will be monitored for oxygenation.

Category

Rehabilitation

Recruitment centers

1

Recruitment center

Name of recruitment center

Baqiyatallah hospital

Full name of responsible person

Marzieh Lak

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Baqiyatallah University of Medical Science, Mollasadra Street, Sheikh Bahaee Street, Tehran, Iran

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Sponsors / Funding sources

1

Sponsor

Name of organization / entity

Bagheiat-allah University of Medical Sciences

Full name of responsible person

Mohammad Hasan Kalantar

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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

Bagheiat-allah 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

Bagheiat-allah University of Medical Sciences

Full name of responsible person

Marzieh Lak

Position

Associate professor

Latest degree

Specialist

Other areas of specialty/work

Anesthesiology

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Person responsible for scientific inquiries

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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

Yes - There is a plan to make this available

Data Dictionary

Yes - There is a plan to make this available

Title and more details about the data/document

All data can be shared after unidentifiable people.

When the data will become available and for how long

Starting 1 month after publication

To whom data/document is available

People working in academic institute

Under which criteria data/document could be used

For meta-analysis

From where data/document is obtainable

by Email

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

By sending a request to the University Vice-Chancellor for Research and obtaining a license

Comments