

Clinical Trial Protocol

Iranian Registry of Clinical Trials

29 Jun 2026

The Effect of Vestibular Stimulation Exercises on Motor Function and Balance in Children with Hearing Impairment

Protocol summary

Study aim

The Effect of Vestibular Stimulation Exercises on Motor Function and Balance in Children with Hearing Impairment

Design

a randomized single-blind clinical trial with intervention and control groups, parallel groups, according to the centralized and computerized study

Settings and conduct

The study is conducted at the preschool school and Ghotbzadeh primary school of Rasht city, as well as single blind design

Participants/Inclusion and exclusion criteria

Hearing-impaired subjects with a hearing loss of about 80 dB are selected. Complete satisfaction and willingness to participate in research. Have no visual impairment

Intervention groups

Intervention group: Vestibular stimulation based on motor exercises. Control group: Do not participate in any exercise program.

Main outcome variables

vestibular, balance

General information

Reason for update

Acronym

IRCT registration information

IRCT registration number: **IRCT20190427043392N1**

Registration date: **2019-06-02, 1398/03/12**

Registration timing: **registered_while_recruiting**

Last update: **2019-06-02, 1398/03/12**

Update count: **0**

Registration date

2019-06-02, 1398/03/12

Registrant information

Name

vahideh sadat mousavi zadeh

Name of organization / entity

Country

Iran (Islamic Republic of)

Phone

+98 34 3411 9618

Email address

v.s.mousavizadeh@gmail.com

Recruitment status

Recruitment complete

Funding source

Expected recruitment start date

2019-05-10, 1398/02/20

Expected recruitment end date

2019-07-11, 1398/04/20

Actual recruitment start date

empty

Actual recruitment end date

empty

Trial completion date

empty

Scientific title

The Effect of Vestibular Stimulation Exercises on Motor Function and Balance in Children with Hearing Impairment

Public title

effect of motor exercises on the balance of children with hearing impairment

Purpose

Supportive

Inclusion/Exclusion criteria

Inclusion criteria:

Hearing-impaired subjects with a hearing loss of about 80 dB are selected. Complete satisfaction and willingness to participate in research. No visual impairment. Children aged 5 to 8 years old

Exclusion criteria:

Beyond the age range of 5 to 8 years. Subjects did not have orthopedic, cognitive, and musculoskeletal disorders. Subjects do not have absolute deafness

Age

From **5 years** old to **8 years** old

Gender

Both

Phase

N/A

Groups that have been masked

No information

Sample size

Target sample size: **24**

Randomization (investigator's opinion)

Randomized

Randomization description

Individuals are randomly assigned randomly into a group or two groups using the random numbers table and they receive the intervention of the same group.

Blinding (investigator's opinion)

Not blinded

Blinding description

Placebo

Not used

Assignment

Parallel

Other design features

Secondary Ids

empty

Ethics committees

1

Ethics committee

Name of ethics committee

Ethics committee of Guilan University of Medical Sciences

Street address

Shahid Siadaty Ave, Namjoo str ,Rasht

City

Rasht

Province

Guilan

Postal code

4188794755

Approval date

2019-03-13, 1397/12/22

Ethics committee reference number

IR. GUMS.REC.1397.479

Health conditions studied

1

Description of health condition studied

hearing loss individual

ICD-10 code

H91

ICD-10 code description

Other and unspecified hearing loss

Primary outcomes

1

Description

balance

Timepoint

measurement of balance tests before and after 6of 6 week intervention

Method of measurement

functional test of balance

2

Description

the timed up and go

Timepoint

subtest of bruninks oseretsky test

Method of measurement

the duration of this test

Secondary outcomes

1

Description

Balance"

Timepoint

Tests for relevant tests after 6 weeks"

Method of measurement

Bronkins - oseretsky test"

2

Description

Motor function

Timepoint

Tests for relevant tests after 6 weeks"

Method of measurement

Bronkins - oseretsky test"

Intervention groups

1

Description

Intervention group: The intervention group carries out a training session on Vestibular stimulation for 6 weeks, every 3 sessions per week.

Category

Rehabilitation

2

Description

Control group: There is no intervention.

Category

N/A

Recruitment centers

1

Recruitment center

Name of recruitment center

Preschool and Deaf Language School, Ghotbzadeh

Full name of responsible person

Pourfathi Hadith

Street address

Preschool school and schoolGotb zadeh elementary,Imam's sister's stepmother, Motehari street

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Province

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4173653137

Phone

+98 13 3335 8501

Email

mghotbzadeh1@gmail.com

Sponsors / Funding sources

1

Sponsor

Name of organization / entity

Rasht University of Medical Sciences

Full name of responsible person

Mirhoseini Seyed Ziaaedin

Street address

5th Kilometer Gazvin Road, Khalij fars Highway,Rasht Down

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Guilan

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4199613776

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+98 13 0338 3369

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szmirhoseini@gmail.com

Grant name

Grant code / Reference number

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

No

Title of funding source

Rasht 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

Rasht University of Medical Sciences

Full name of responsible person

Mousavi zadeh Vahideh sadat

Position

Msc Student

Latest degree

Bachelor

Other areas of specialty/work

Sport injory and corrective exercise

Street address

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Contact

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Rasht University of Medical Sciences

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Latest degree

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Person responsible for updating data

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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 variable data measured after unidentifiable
individuals can be shared

When the data will become available and for how long

Starting in January 2020

To whom data/document is available

Only for researchers working in academic and scientific
institutions

Under which criteria data/document could be used

Scientific use in improving and comparing with other
research is allowed

From where data/document is obtainable

E-mail address for correspondence

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

Applicants should send the reason for using the
documentation to an E-mail.

Comments