

# Clinical Trial Protocol

## Iranian Registry of Clinical Trials

09 Jun 2026

### Effect of co-enzyme Q10 versus placebo on treatment of tinnitus due to presbycusis: a triple-blind randomized clinical trial

#### Protocol summary

##### Study aim

To assess the effect of co-enzyme Q10 versus placebo on treatment of tinnitus due to presbycusis

##### Design

This is a triple-blind randomized clinical trial, phase II, in which 50 eligible patients will be randomly assigned to the intervention and control groups

##### Settings and conduct

The eligible patients with tinnitus due to presbycusis referring to the Besat Hospital in Hamadan city during the study period will be enrolled in the trial and will be randomly assigned to the intervention and control groups through the drawing of lots. This trial will be triple-blinded so that neither patients nor the physician examining the patients and nor the data analyzer will be aware of the intervention.

##### Participants/Inclusion and exclusion criteria

Inclusion criteria: Age of 50 to 90 years, Tinnitus for over a year, Bilateral neural sensorineural hearing loss  
Exclusion criteria: Psychosomatic disorder, Consumption of calcium, vitamin, and magnesium in the past two months, Tinnitus due to drug side effects, Atypical tinnitus, Meniere's disease, Conductive hearing loss

##### Intervention groups

Intervention group: Neurotriptyline tablet 25 mg (manufactured by Darupakhsh Pharmaceutical Co.) daily plus co-enzyme Q10 tablet 100 mg (manufactured by Health Aid Co.) daily for 6 weeks  
Control group: Neurotriptyline tablet 25 mg (manufactured by Darupakhsh Pharmaceutical Co.) daily plus placebo tablet 100 mg daily for 6 weeks

##### Main outcome variables

Primary outcome: The severity of tinnitus, sleep disorder, quality of life

#### General information

##### Reason for update

##### Acronym

#### IRCT registration information

IRCT registration number: **IRCT20120215009014N339**

Registration date: **2020-02-07, 1398/11/18**

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

Last update: **2020-02-07, 1398/11/18**

Update count: **0**

#### Registration date

2020-02-07, 1398/11/18

#### Registrant information

##### Name

Jalal Poorolajal

##### Name of organization / entity

Department of Epidemiology & Biostatistics Hamadan University of Medical Sciences

##### Country

Iran (Islamic Republic of)

##### Phone

+98 81 1838 0090

##### Email address

poorolajal@umsha.ac.ir

#### Recruitment status

**Recruitment complete**

#### Funding source

#### Expected recruitment start date

2019-09-23, 1398/07/01

#### Expected recruitment end date

2020-02-19, 1398/11/30

#### Actual recruitment start date

empty

#### Actual recruitment end date

empty

#### Trial completion date

empty

#### Scientific title

Effect of co-enzyme Q10 versus placebo on treatment of tinnitus due to presbycusis: a triple-blind randomized

clinical trial

## Public title

Effect of co-enzyme Q10 versus placebo on treatment of tinnitus due to presbycusis

## Purpose

Treatment

## Inclusion/Exclusion criteria

### Inclusion criteria:

Age of 50 to 90 years, Tinnitus for over a year, Bilateral neural sensorineural hearing loss

### Exclusion criteria:

Psychosomatic disorder, Consumption of calcium, vitamin and magnesium in the past two months, Tinnitus due to drug side effects, Atypical tinnitus, Meniere's disease, Conductive hearing loss

## Age

From **50 years** old to **90 years** old

## Gender

Both

## Phase

2

## Groups that have been masked

- Participant
- Care provider
- Investigator
- Outcome assessor
- Data analyser
- Data and Safety Monitoring Board

## Sample size

Target sample size: **50**

## Randomization (investigator's opinion)

Randomized

## Randomization description

Random assignment of the patients to the intervention and control groups through the drawing of lots. To do this, we prepare two sheets and write "intervention" on one sheet and "control" on another. Then, by referring each patient, one of the sheets will be randomly taken and the patient will be assigned to the intervention or

## Blinding (investigator's opinion)

Triple blinded

## Blinding description

The drugs will be given in coded envelopes. The shape of the medications and placebos will be perfectly the same. Therefore, patients will be unaware of the type of intervention. The physician who will examine the patients will not be aware of the intervention. The analyzer will be unaware of the type of interventions. Thus, the trial will be run as triple-blind

## Placebo

Used

## Assignment

Parallel

## Other design features

## Secondary Ids

empty

## Ethics committees

### 1

#### Ethics committee

##### Name of ethics committee

Ethics Committee of Hamadan University of Medical Sciences

##### Street address

Vice-chancellor for Research and Technology, Hamadan University of Medical Sciences, Shahid Fahmideh Ave

##### City

Hamadan

##### Province

Hamadan

##### Postal code

6517838695

#### Approval date

2019-07-06, 1398/04/15

#### Ethics committee reference number

IR.UMSHA.REC.1398.301

## Health conditions studied

### 1

#### Description of health condition studied

Tinnitus

#### ICD-10 code

H93.1

#### ICD-10 code description

Tinnitus

## Primary outcomes

### 1

#### Description

Severity of tinnitus

#### Timepoint

Before the intervention and 6 months later

#### Method of measurement

Using the Tinnitus Handicap Inventory (THI)

### 2

#### Description

Sleep disorder

#### Timepoint

Before the intervention and 6 months later

#### Method of measurement

Using the Petersburg questionnaire

### 3

#### Description

Quality of life

#### Timepoint

Before the intervention and 6 months later

#### Method of measurement

Using the SF-36 questionnaire

## Secondary outcomes

empty

## Intervention groups

1

### Description

Intervention group: Neurotriptyline tablet 25 mg (manufactured by Darupakhsh Pharmaceutical Co.) daily plus co-enzyme Q10 tablet 100 mg (manufactured by Health Aid Co.) daily for 6 weeks

### Category

Treatment - Drugs

2

### Description

Control group: Neurotriptyline tablet 25 mg (manufactured by Darupakhsh Pharmaceutical Co.) daily plus placebo tablet 100 mg daily for 6 weeks

### Category

Placebo

## Recruitment centers

1

### Recruitment center

#### Name of recruitment center

Besat Hospital in Hamadan city

#### Full name of responsible person

Dr Zahra Sadeghi

#### Street address

Besat Hospital, Shahed Square

#### City

Hamadan

#### Province

Hamadan

#### Postal code

6517838695

#### Phone

+98 81 3364 0030

#### Email

sazcsaz66@gmail.com

## Sponsors / Funding sources

1

### Sponsor

#### Name of organization / entity

Hamedan University of Medical Sciences

#### Full name of responsible person

Dr. Saeid Bashirian

#### Street address

Hamadan University of Medical Sciences, Shahid Fahmideh Ave

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#### Postal code

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#### Phone

+98 81 3838 0717

#### Email

info.research@umsha.ac.ir

#### Grant name

#### Grant code / Reference number

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

Yes

#### Title of funding source

Hamedan 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

Hamedan University of Medical Sciences

#### Full name of responsible person

Dr Zahra Sadeghi

#### Position

Resident of ENT

#### Latest degree

Medical doctor

#### Other areas of specialty/work

Ear, Nose, and Throat

#### Street address

School of Medicine, Hamadan University of Medical Sciences, Shahid Fahmideh Ave.

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6517838695

#### Phone

+98 81 3838 0572

#### Email

sazcsaz66@gmail.com

## Person responsible for scientific inquiries

### Contact

#### Name of organization / entity

Hamedan University of Medical Sciences

#### Full name of responsible person

Dr. Rooholah Abbasi

#### Position

Otolaryngologist

**Latest degree**

Medical doctor

**Other areas of specialty/work**

Ear, Nose, and Throat

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Ruholah@abasihatmil.com

## Person responsible for updating data

**Contact**

**Name of organization / entity**

Hamedan University of Medical Sciences

**Full name of responsible person**

Dr. Jalal Poorolajal

**Position**

Professor of Epidemiology

**Latest degree**

Ph.D.

**Other areas of specialty/work**

Epidemiology

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**Email**

poorolajal@umsha.ac.ir

## 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