

# Clinical Trial Protocol

## Iranian Registry of Clinical Trials

28 May 2026

### Study of adding *Boswellia serrata* resin to therapy of patients with mild to moderate Alzheimer's disease on cognitive status compared with placebo

#### Protocol summary

##### Study aim

Evaluation of the effect of *Olibanum* resin on improvement of mild to moderate Alzheimer's disease

##### Design

30 sample size RCT in tow parallel group placebo&Control, Double-Blind, Block Randomization using the website [www.Randomization.com](http://www.Randomization.com)

##### Settings and conduct

Alzheimer's patients referring to clinic of Avicenna's hospital in Mashhad if they have inclusion criteria after the signature of the informed consent form by legal guardians are randomly and double blind (such as Researchers, clinical caregivers, patients, outcome evaluator, and data analyzer) divided into two groups of 15 treatment and placebo; after Receiving capsules for 12 weeks; Mental status of both groups is studying with MMSE and CDR tests and compared to week zero Using SPSS.16 software.

##### Participants/Inclusion and exclusion criteria

Inclusion criteria: Patients with mild to moderate Alzheimer's disease based on the score of the MMSE test= 10-23 Exclusion criteria: Diagnosis of Alzheimer's disease less than 6 months before entering the study, Using cholinesterase inhibitors and / or memantine less than 90 days before entering the study or discontinuing them less than 90 days before entering the study, Use of other drugs and substances that affect cognitive status less than 60 days before entering the study or discontinued less than 90 days before entering the study, Vascular dementia, thyroid disease, vitamin B12 deficiency, anemia, Lewy body dementia, liver function disorders, brain tumors, ischemic CVA

##### Intervention groups

Intervention group: A capsule containing 750 mg of *Olibanum* resin is taken orally three times per day for 12 weeks. Control group: The placebo is used like *Olibanum* in the form of capsules are apparently similar to the

study drug , which contains 500 mg of starch powder. Capsules are provided by the Pharmacy Department of Herbal Drugs at Imam Reza Pharmacy in Mashhad

##### Main outcome variables

Cognitive status

#### General information

##### Reason for update

##### Acronym

##### IRCT registration information

IRCT registration number: **IRCT20161219031463N2**

Registration date: **2019-03-09, 1397/12/18**

Registration timing: **retrospective**

Last update: **2019-03-09, 1397/12/18**

Update count: **0**

##### Registration date

2019-03-09, 1397/12/18

##### Registrant information

##### Name

Hamid Reza Sadeghnia

##### Name of organization / entity

##### Country

Iran (Islamic Republic of)

##### Phone

+98 51 3882 8566

##### Email address

sadeghniahr@mums.ac.ir

##### Recruitment status

**Recruitment complete**

##### Funding source

##### Expected recruitment start date

2018-10-09, 1397/07/17

##### Expected recruitment end date

2018-12-22, 1397/10/01

**Actual recruitment start date**

empty

**Actual recruitment end date**

empty

**Trial completion date**

empty

**Scientific title**

Study of adding Boswellia serrata resin to therapy of patients with mild to moderate Alzheimer's disease on cognitive status compared with placebo

**Public title**

Improving effects of olibanum on learning and memory

**Purpose**

Treatment

**Inclusion/Exclusion criteria****Inclusion criteria:**

Patients with mild to moderate Alzheimer's disease based on the score of the MMSE test between 10-23

**Exclusion criteria:**

Diagnosis of Alzheimer's disease less than 6 months before entering the study Using cholinesterase inhibitors and / or memantine less than 90 days before entering the study or discontinuing them less than 90 days before entering the study Use of other drugs and substances such as ginkgo biloba or vitamin E that affect cognitive status less than 60 days before entering the study or discontinued less than 90 days before entering the study. Vascular dementia, thyroid disease, vitamin B12 deficiency, anemia, Lewy body dementia, liver function disorders, brain tumors, ischemic cerebrovascular accidents

**Age**

From **55 years** old to **82 years** old

**Gender**

Both

**Phase**

2

**Groups that have been masked**

- Participant
- Care provider
- Investigator
- Outcome assessor
- Data analyzer

**Sample size**

Target sample size: **30**

**Randomization (investigator's opinion)**

Randomized

**Randomization description**

Randomization was done individually, block and using the website www.Randomization.com with 6 subjects per block. This was done by an independent researcher and was given to the capsule deliverer by naming the unknown letters that the person did not know the original name of the capsules named in letters.

**Blinding (investigator's opinion)**

Double blinded

**Blinding description**

Groups as participants, clinical care givers, outcome evaluators and data analyzer are not aware of the

presence of samples in the control group of the placebo or the study group. So that the duty of dividing the patients into two groups was performed by the person who was not dependent on the study, and the distribution of the drug and placebo was done by the same person, and until the end of the study, information about the grouping of patients was reserved by this person

**Placebo**

Used

**Assignment**

Parallel

**Other design features****Secondary Ids**

empty

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

The Ethics Committee of Mashhad University of Medical Sciences

**Street address**

Ghershi building, Daneshgah Ave.

**City**

Mashhad

**Province**

Razavi Khorasan

**Postal code**

91388-13944

**Approval date**

2016-12-18, 1395/09/28

**Ethics committee reference number**

IR.HUHS.REC.1395.438

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

Alzheimer's disease

**ICD-10 code**

F00

**ICD-10 code description**

Dementia in Alzheimer disease

**Primary outcomes****1****Description**

Cognitive status

**Timepoint**

Study of cognitive status at the beginning of the study (before the intervention) and 12 weeks after the start of use of capsule containing Olibanum resin or placebo

**Method of measurement**

Using MMSE and CDR tests

## Secondary outcomes

empty

## Intervention groups

### 1

#### Description

Intervention group: A capsule containing 750 mg of Olibuma resin is taken orally three times per day for 12 weeks. Capsules are provided by the Pharmacy Department of Herbal Drugs at Imam Reza Pharmacy in Mashhad

#### Category

Treatment - Drugs

### 2

#### Description

Control group: The placebo is used in the form of capsules are apparently similar to the study drug , which contains 500 mg of starch powder three times a day for 12 weeks orally. Capsules are provided by the pharmacy department of Herbal Drugs at Imam Reza Pharmacy in Mashhad

#### Category

Placebo

## Recruitment centers

### 1

#### Recruitment center

##### Name of recruitment center

Ebne Sina hospital

##### Full name of responsible person

Ali Manteghi

##### Street address

Next to Astan Ghods Razavi's transition, Horr Ameli Blv.

##### City

Mashhad

##### Province

Razavi Khorasan

##### Postal code

919583134

##### Phone

+98 51 3711 2701

##### Fax

+98 51 3711 2545

##### Email

ManteghiA@mums.ac.ir

##### Web page address

<http://sina.mums.ac.ir>

## Sponsors / Funding sources

### 1

#### Sponsor

Name of organization / entity

Mashhad University of Medical Sciences

#### Full name of responsible person

Mohsen Tafaghodi

#### Street address

Ghershi building, Daneshgah Ave.

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#### Phone

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#### Fax

+98 51 3882 3251

#### Email

[tafaghodim@mums.ac.ir](mailto:tafaghodim@mums.ac.ir)

#### Web page address

<http://v-research.mums.ac.ir/index.php/moaven/tafaghodim>

#### Grant name

940998

#### Grant code / Reference number

940998

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

Yes

#### Title of funding source

Mashhad 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

Mashhad University of Medical Sciences

##### Full name of responsible person

Hamid Reza Sadeghnia

##### Position

Associate Professor

##### Latest degree

Ph.D.

##### Other areas of specialty/work

Medical Pharmacy

##### Street address

Pharmacology Department, Faculty of Medicine, Inside campus college, Azadi square

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##### Province

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##### Postal code

91388-13944

**Phone**

+98 51 3800 2257

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+98 51 3882 8567

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Sadeghniahr@mums.ac.ir

**Web page address**

<http://pharmacodept.mums.ac.ir>

## Person responsible for scientific inquiries

**Contact**

**Name of organization / entity**

Mashhad University of Medical Sciences

**Full name of responsible person**

Hami Reza Sadeghnia

**Position**

Associate Professor

**Latest degree**

Ph.D.

**Other areas of specialty/work**

Medical Pharmacy

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## Person responsible for updating data

**Contact**

**Name of organization / entity**

Mashhad University of Medical Sciences

**Full name of responsible person**

Hamid Reza Sadeghnia

**Position**

Associate Professor

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**Web page address**

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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**

No - There is not 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**

Demographic data, MMSE and CDR tests' scores

**When the data will become available and for how long**

Start the access period 6 months after Releasing the results

**To whom data/document is available**

Researchers working in academic and scientific institutions

**Under which criteria data/document could be used**

For use in complementary studies including repeat study with higher sample size

**From where data/document is obtainable**

Dr Hamid Reza Sadeghnia, Mashhad Faculty of Medicine, Department of Pharmacology Email: Sadeghniahr@mums.ac.ir Fax: +98 51 3882 8567 Address: Pharmacology Department, Faculty of Medicine, Inside campus college, Azadi square Postal code: 91388-13944

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

Submit a formal and written request from the Research, Academic and Scientific Institute at the address above

**Comments**

No comments