

Clinical Trial Protocol

Iranian Registry of Clinical Trials

07 Jul 2026

Comparing the effect of violet almond oil with almond oil and placebo in quality of sleep in patients with chronic insomnia

Protocol summary

Summary

Primary insomnia patients were detected by a neurologist based on the diagnostic criteria in DSM-V. The included patients were those whose illness had lasted at least three months and were willing to participate in a research project. The patients were referred to the Clinic of Traditional and Complementary Medicine in Mashhad University of Medical Sciences. After obtaining informed consent, they were randomly divided into three groups: A, B, C. The three groups received three drops of almond oil or violet-almond oil or placebo on each nostril every night. Since the treatment of chronic insomnia cares about sleep hygiene, due to moral considerations and lack of deprivation of treatment in the placebo group, sleep hygiene instructions were presented to all the three groups based on the guidelines of the American Association sleep (AASM). Before and after 30 days of treatment, Pittsburgh Sleep Quality Index, Insomnia Severity, was completed by all patients.

General information

Acronym

IRCT registration information

IRCT registration number: **IRCT2015120125335N1**

Registration date: **2016-02-19, 1394/11/30**

Registration timing: **registered_while_recruiting**

Last update:

Update count: **0**

Registration date

2016-02-19, 1394/11/30

Registrant information

Name

Zohre Feyzabadi

Name of organization / entity

Faculty of Traditional and Complementary Medicine,
Mashhad University of Medical Sciences

Country

Iran (Islamic Republic of)

Phone

+98 51 3855 2188

Email address

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Recruitment status

Recruitment complete

Funding source

Mashhad University of Medical Sciences

Expected recruitment start date

2016-01-21, 1394/11/01

Expected recruitment end date

2016-05-21, 1395/03/01

Actual recruitment start date

empty

Actual recruitment end date

empty

Trial completion date

empty

Scientific title

Comparing the effect of violet almond oil with almond oil and placebo in quality of sleep in patients with chronic insomnia

Public title

Comparing the effect of violet almond oil with almond oil and placebo in chronic insomnia

Purpose

Treatment

Inclusion/Exclusion criteria

Inclusion criteria: Willingness to cooperate; Having the pre-defined conditions of insomnia according to DSM-V-TR; Suffering from insomnia for at least 3 months; Being between 18 and 60 years old; Lack of physical diseases such as lung diseases, allergic rhinitis, cardiovascular, infectious diseases, cancer; Lack of mental illnesses such

as depression, substance and drug abuse; Lack of neurological diseases such as Parkinson's and Alzheimer's disease; lack of sleep disorders like sleep apnea; lack of using drug and non-drug treatments for insomnia. Exclusion criteria: Pregnant and lactating women; History of allergy to medicinal herbs; Side effects of drugs; Irregular use of medication or treatment discontinuation; unwillingness to continue cooperation for any reason.

Age

From **18 years** old to **60 years** old

Gender

Both

Phase

N/A

Groups that have been masked

No information

Sample size

Target sample size: **75**

Randomization (investigator's opinion)

Randomized

Randomization description

Blinding (investigator's opinion)

Double blinded

Blinding description

Placebo

Used

Assignment

Parallel

Other design features

Secondary Ids

empty

Ethics committees

1

Ethics committee

Name of ethics committee

Mashhad University of Medical Sciences

Street address

Building Qureshi, Daneshgah Ave.

City

Mashhad

Postal code

Approval date

2015-12-09, 1394/09/18

Ethics committee reference number

IR.mums.fm.REC.1394.461

Health conditions studied

1

Description of health condition studied

chronic insomnia

ICD-10 code

F51.0

ICD-10 code description

Nonorganic insomnia

Primary outcomes

1

Description

sleep

Timepoint

Before the intervention and one month after intervention

Method of measurement

Pittsburgh questionnaire, Insomnia Severity Index questionnaire

Secondary outcomes

1

Description

Side effects

Timepoint

The end of study

Method of measurement

Questionnaire

Intervention groups

1

Description

In the first group of intervention, patients receive 3 drops of almond violet oil in each nostril every night before sleep for a duration of one month. Each drop contains 33 mg of the drug.

Category

Treatment - Drugs

2

Description

In the placebo group, patients receive 3 drops of placebo in each nostril every night before sleep for a duration of one month. Each drop contains 33 mg of the placebo.

Category

Placebo

3

Description

In the second group of intervention, patients receive 3 drops of sweet almond oil in each nostril every night before sleep for a duration of one month. Each drop contains 33 mg of the drug.

Category

Treatment - Drugs

Recruitment centers

1

Recruitment center

Name of recruitment center

Traditional Medicine Clinic

Full name of responsible person

Zohre Feyzabadi

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East Razi Ave.**City**

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Web page address**Person responsible for scientific inquiries****Contact****Name of organization / entity**Faculty of Traditional and Complementary Medicine,
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Position

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Web page address**Sponsors / Funding sources****1****Sponsor****Name of organization / entity**

Mashhad University of Medical Sciences

Full name of responsible person

Mohsen Tafaghodi

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City

Mashhad

Grant name**Grant code / Reference number****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*empty***Domestic or foreign origin***empty***Category of foreign source of funding***empty***Country of origin****Type of organization providing the funding***empty***Person responsible for general inquiries****Contact****Name of organization / entity**Faculty of Traditional and Complementary Medicine,
Mashhad University of Medical Sciences**Full name of responsible person**

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Web page address**Sharing plan****Deidentified Individual Participant Data Set (IPD)***empty***Study Protocol***empty***Statistical Analysis Plan***empty***Informed Consent Form**

empty
Clinical Study Report
empty
Analytic Code

empty
Data Dictionary
empty