

Clinical Trial Protocol

Iranian Registry of Clinical Trials

12 Jul 2026

Comparison the effects of memantine and sodium valproate in migraine pain prophylaxis

Protocol summary

Study aim

Comparison the effects of memantine and sodium valproate in migraine pain prophylaxis

Design

The clinical trial in phase 3 has two groups, community-based with parallel groups, double-blind and pseudo randomization methods

Settings and conduct

Migraine patients referring to neurology clinics of Ardabil University of Medical Sciences will be reviewed. This study will be done as double-blinded form (blindness of patients, researcher and physicians). Patients will be divided into two groups of 40 by pseudo randomization method. We will use Memantine for a group and sodium valproate for another group for 12 weeks. After taking their medications, patients do not receive any medication for 4 weeks. Then we will give memantine to sodium valproate group and sodium valproate to memantine group for another 12 weeks. Patients will be visited every four weeks by their neurologist according to a pre-prepared questionnaire.

Participants/Inclusion and exclusion criteria

Inclusion criteria: Migraine history, more than 3 headache attacks per month, prophylactic treatment failure with one or more migraine prevention drugs, aged 16 to 65 years, spacing more than 48 hours between attacks
Exclusion criteria: Other headaches, use of other migraine prevention medications, drug dependence on drugs or alcohol, renal failure, excessive use of acute migraine attack drugs

Intervention groups

Group 1: These patients will receive 10mg of memantine coated tablets per day and will be continued for 12 weeks. Group 2: These patients will receive 400mg of sodium valproate coated tablets daily and will continue for 12 weeks. Then patients will not take any drug for 4 weeks and then will be given sodium valproate to patients in group 1 and memantine to patients in group 2 for 12 weeks

Main outcome variables

Headache score by MIDAS score

General information

Reason for update

Acronym

IRCT registration information

IRCT registration number: **IRCT20180624040229N1**

Registration date: **2019-01-02, 1397/10/12**

Registration timing: **retrospective**

Last update: **2019-01-02, 1397/10/12**

Update count: **0**

Registration date

2019-01-02, 1397/10/12

Registrant information

Name

Rogayye Dargahi

Name of organization / entity

Country

Iran (Islamic Republic of)

Phone

+98 45 3323 0739

Email address

arumail@arums.ac.ir

Recruitment status

Recruitment complete

Funding source

Expected recruitment start date

2018-01-21, 1396/11/01

Expected recruitment end date

2018-10-12, 1397/07/20

Actual recruitment start date

2018-01-21, 1396/11/01

Actual recruitment end date

2018-10-12, 1397/07/20

Trial completion date

2018-10-12, 1397/07/20

Scientific title

Comparison the effects of memantine and sodium valproate in migraine pain prophylaxis

Public title

Comparison the effects of memantine and sodium valproate in migraine pain prophylaxis

Purpose

Prevention

Inclusion/Exclusion criteria**Inclusion criteria:**

A history of migraines based on the criteria of the international community of headache. The number of headache attacks is more than three attacks per month. Failure of prophylactic treatment with one or more migraine prevention drugs. Any headache attack with more than 48 hours of analgesia should be distinguished from other attacks Filling out the consent form of the Clinical trial age 16 to 65

Exclusion criteria:

Other headache Use of other migraine prevention medications Drug dependence on drugs or alcohol Renal dysfunction Excessive use of acute migraine attack drugs (more than 8 days in the month of receiving ergots, NSAIDs or tryptans) Sensitization to memantine or sodium valproate or their compounds. Acute resistant migraine Severe psychiatric disorders (bipolar disorder, mental disorder, severe depressive disorder, obsessive-compulsive disorder, or functional disorder, general anxiety disorder, Somatization) Neurodegenerative disorders malignancy , pain disorder, severe infection Pregnant or Breastfeeding women

Age

From **16 years** old to **65 years** old

Gender

Both

Phase

3

Groups that have been masked

- Participant
- Care provider
- Investigator

Sample size

Target sample size: **80**

Actual sample size reached: **80**

Randomization (investigator's opinion)

Not randomized

Randomization description**Blinding (investigator's opinion)**

Double blinded

Blinding description

The way of participant blindness is that he will receive and use a certain number of drugs in an unnamed and labeled envelope. The way of physicians blindness is that after he has been diagnosed migraine to his patient and notice his secretary, randomly given one of the envelopes to the patient. The way of researcher's blindness is that he will not be present in the filling

envelopes and selecting them.

Placebo

Not used

Assignment

Crossover

Other design features**Secondary Ids**

empty

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

Ethics committee of Ardabil University of Medical Sciences

Street address

No. 1, Moallem street,

City

Ardabil

Province

Ardabil

Postal code

5613843793

Approval date

2017-10-23, 1396/08/01

Ethics committee reference number

IR.ARUMS.REC.1396.80

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

Migraine

ICD-10 code

G43.0

ICD-10 code description

Migraine without aura

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

Headache score by MIDAS score

Timepoint

At the beginning , 3 month later, 6 month later

Method of measurement

MIDAS score

Secondary outcomes

empty

Intervention groups

1

Description

First intervention group: Memantine, 5 mg oral tablet (Sobhan company), twice a day for 12 weeks. Then 4 weeks without taking any medication. Then take 200 mg of sodium valproate (Raha company), twice a day for 12 weeks

Category

Treatment - Drugs

2

Description

Second Intervention group: Sodium valproate, 200 mg oral tablet (Raha company), twice a day for 12 weeks. Then 4 weeks without taking any medication. Then take 5 mg of Memantine (Sobhan company), twice a day for 12 weeks

Category

Treatment - Drugs

Recruitment centers

1

Recruitment center

Name of recruitment center

Alavi Hospital

Full name of responsible person

Meysam Fouladi

Street address

Royan clinic, Pasdaran St

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Ardabil

Postal code

5613843793

Phone

+98 45 3323 0739

Email

Reseteam.2020@gmail.com

Sponsors / Funding sources

1

Sponsor

Name of organization / entity

Ardabil University of Medical Sciences

Full name of responsible person

Parvaneh Naftchi

Street address

Ardabil university of medical sciences, Daneshgah St

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Province

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Postal code

5618985991

Phone

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Email

info@arums.ac.ir

Web page address

Grant name

Grant code / Reference number

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

Yes

Title of funding source

Ardabil 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

Ardabil University of Medical Sciences

Full name of responsible person

Vahid Abbasi

Position

Associate professor

Latest degree

Specialist

Other areas of specialty/work

Neurology

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Person responsible for scientific inquiries

Contact

Name of organization / entity

Ardabil University of Medical Sciences

Full name of responsible person

Vahid Abbasi

Position

Associate professor

Latest degree

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Person responsible for updating data

Contact

Name of organization / entity
Ardabil University of Medical Sciences
Full name of responsible person
Vahid Abbasi
Position
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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

No - There is not 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

Undecided - It is not yet known if there will be a plan to make this available

Title and more details about the data/document

After unidentified people, demographic data and MIDAS score will be used to evaluate the headache.

When the data will become available and for how long

Starting the access from 2020

To whom data/document is available

Access will be free for everyone

Under which criteria data/document could be used

Statistical analyzes are allowed on data

From where data/document is obtainable

Will be contact by Dr Abbasi`s email

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

It takes a total of one month

Comments

No other information is required