

# Clinical Trial Protocol

## Iranian Registry of Clinical Trials

24 Jun 2026

### Randomized placebo controlled clinical trial of the efficacy of Gabapentin on flushing in menopausal women

#### Protocol summary

##### Summary

This randomized single blind placebo controlled clinical trial will be conducted on sixty menopausal women, who elapse from their last menstruation at least one year and FSH level more than 40 IU/L. Exclusion criteria are: consumption of any drugs for flushing, having systemic diseases, happening the Gabapentin complications noncompliance. The aim of this study is to evaluate the efficacy of Gabapentin on the severity of flushing. The patients will randomly assign into intervention and control groups. In intervention group patients will take Gabapentin capsules (300mg), three times a day, for three months. The control group will receive placebo capsules (consist of starch) three times a day for three months. Primary outcome measure is the severity of flushing measured by visual analog scale at the beginning, first, second and third months after drug consumption. The results will be compared in study groups.

#### General information

##### Acronym

##### IRCT registration information

IRCT registration number: **IRCT201103011107N4**  
Registration date: **2011-05-05, 1390/02/15**  
Registration timing: **retrospective**

Last update:

Update count: **0**

##### Registration date

2011-05-05, 1390/02/15

##### Registrant information

##### Name

Tajea Jadery

##### Name of organization / entity

Jundi shapur university

##### Country

Iran (Islamic Republic of)

##### Phone

-

##### Email address

research@ajums.ac.ir

##### Recruitment status

**Recruitment complete**

##### Funding source

Jundshapour University of Medical Sciences

##### Expected recruitment start date

2010-06-22, 1389/04/01

##### Expected recruitment end date

2010-10-23, 1389/08/01

##### Actual recruitment start date

empty

##### Actual recruitment end date

empty

##### Trial completion date

empty

##### Scientific title

Randomized placebo controlled clinical trial of the efficacy of Gabapentin on flushing in menopausal women

##### Public title

Efficacy of Gabapentin on flushing in menopausal women

##### Purpose

Treatment

##### Inclusion/Exclusion criteria

Inclusion criteria : Elapsing one year of last menstrual phase(LMP), Follicle-stimulating hormone (FSH) more than 40IU/L and not using any drugs that affect on hot flush symptoms. Exclusion criteria : happening Gabapentin side effects, noncompliance and coexistence of systemic disease

##### Age

From **40 years** old to **55 years** old

##### Gender

Female

## Phase

2-3

## Groups that have been masked

No information

## Sample size

Target sample size: 60

## Randomization (investigator's opinion)

Randomized

## Randomization description

## Blinding (investigator's opinion)

Single blinded

## Blinding description

## Placebo

Used

## Assignment

Parallel

## Other design features

## Secondary Ids

empty

## Ethics committees

### 1

#### Ethics committee

##### Name of ethics committee

Ahwaz Jondishapur University of Medical Sciences

##### Street address

Ahwaz

##### City

Golestan Bolvar,Ahwaz

##### Postal code

#### Approval date

2010-06-07, 1389/03/17

#### Ethics committee reference number

2380

## Health conditions studied

### 1

#### Description of health condition studied

Flushing

#### ICD-10 code

N95.1

#### ICD-10 code description

Menopausal and female climacteric states

## Primary outcomes

### 1

#### Description

Flushing severity

#### Timepoint

Before intervention, 1 month after intervention, 2months later and 3 months later after intervention

#### Method of measurement

Visual Analoge Scale

## Secondary outcomes

### 1

#### Description

Flushing duration

#### Timepoint

Before intervention, 1 month after intervention, 2months later and 3 months later after intervention

#### Method of measurement

Patients notation

## Intervention groups

### 1

#### Description

Intervention group: Gabapentin Capsule 300 mg- TDS- during 3 months

#### Category

Treatment - Drugs

### 2

#### Description

Control group: Placebo as capsules, three times a day for 3 months

#### Category

Placebo

## Recruitment centers

### 1

#### Recruitment center

##### Name of recruitment center

Emam Khomeini Hospital

##### Full name of responsible person

Solmaz Natanj

##### Street address

Ahwaz Emam Khomeini Hospital

##### City

Ahwaz

## Sponsors / Funding sources

### 1

#### Sponsor

##### Name of organization / entity

Jundishapour Medical University

##### Full name of responsible person

Dr Mostafa Fegghi,Vice-Chancellor For Research

##### Street address

Vice-Chancellor For Research, Ahwaz Jundishapour University of Medical Science

##### City

Ahwaz

#### Grant name

#### Grant code / Reference number

#### Is the source of funding the same sponsor

**organization/entity?**

Yes

**Title of funding source**

Jundishapour Medical University

**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**

Ahvaz Jundishapour Medical University

**Full name of responsible person**

Dr. Solmaz Natanj

**Position**

Obstetrics &amp; Gynecology Resident

**Other areas of specialty/work****Street address**

Ahvaz Emam Khomeini Hospital

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Solmaznatanj@Yahoo.com

**Web page address****Person responsible for scientific inquiries****Contact****Name of organization / entity**

Ahvaz Jundishapour Medical University

**Full name of responsible person**

Dr. Najmieh Saadati

**Position**

Assistant Professor Of Obstetrics &amp; Gynecology

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Ahvaz Jundishapour Medical University

**Full name of responsible person**

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**Position**

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**Other areas of specialty/work****Street address**

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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*