

# Clinical Trial Protocol

## Iranian Registry of Clinical Trials

12 Jun 2026

### Evaluate the effect of Vitagnus on premenstrual syndrome in female students of Shahid Chamran dormitory

#### Protocol summary

##### Summary

This study aimed to determine the effect of Vitagnus on premenstrual syndrome in female students of Shahid Chamran dorm, and it was conducted as a double-blind clinical trial. Inclusion criterion: Female students suffering PMS with regular menstrual cycle and exclusion criteria: current psychotherapy, breast feeding/pregnancy, taking sexual hormones, having stress in the past three months (e.g. death, marriage or surgery of close relatives). In this study, after completing the questionnaire twice during three consecutive menstrual cycles, 72, daughter of Shahid Chamran dormitory students who 25\_18 years of age, suffering from moderate to severe premenstrual syndrome and the conditions for inclusion were selected and then randomly divided into experimental and control groups were 36. Case group was prescribed with 40 drops of Vitagnus daily for three menstrual cycles, while the control group was given placebo.

#### General information

##### Acronym

##### IRCT registration information

IRCT registration number: **IRCT2014102219632N1**  
Registration date: **2015-02-04, 1393/11/15**  
Registration timing: **prospective**

Last update:

Update count: **0**

##### Registration date

2015-02-04, 1393/11/15

##### Registrant information

##### Name

Parvaneh Mousavi

##### Name of organization / entity

Ahvaz University Of Medical Science

##### Country

Iran (Islamic Republic of)

##### Phone

+98 61 3554 4902

##### Email address

mousavi-p@ajums.ac.ir

##### Recruitment status

**Recruitment complete**

##### Funding source

Reseach Deputy of Ahvaz University of Medical Sciences

##### Expected recruitment start date

2015-03-20, 1393/12/29

##### Expected recruitment end date

2015-03-20, 1393/12/29

##### Actual recruitment start date

empty

##### Actual recruitment end date

empty

##### Trial completion date

empty

##### Scientific title

Evaluate the effect of Vitagnus on premenstrual syndrome in female students of Shahid Chamran dormitory

##### Public title

Vitagnus effect on premenstrual symptoms

##### Purpose

Treatment

##### Inclusion/Exclusion criteria

Inclusion criteria: students with premenstrual syndrome with regular menstrual cycle. exclusion criteria: Psychotherapy with pregnancy or breastfeeding; use of hormones; stress in the recent quarter such as marriage or the death of someone close surgery; patients taking certain medications such as dopamine antagonists used

##### Age

From **18 years** old to **25 years** old

##### Gender

Female

## Phase

3

## Groups that have been masked

No information

## Sample size

Target sample size: 72

## 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

Ahvaz University of Medical Sciences

#### Street address

Ahvaz University of Medical Sciences, Esfand Street,  
Golestan

#### City

Ahvaz

#### Postal code

### Approval date

2003-01-21, 1381/11/01

### Ethics committee reference number

278

## Health conditions studied

1

### Description of health condition studied

Premenstrual Syndrome

### ICD-10 code

N94.3

### ICD-10 code description

Unspecified condition associated with female genital  
organs and menstrual cycle

## Primary outcomes

1

### Description

the severity of the symptoms of premenstrual syndrome

### Timepoint

one: two Three months after intervention

## Method of measurement

Recorded daily symptoms of PMS

## Secondary outcomes

1

### Description

Side effects include itching, dizziness, stomach pain,  
diarrhea

### Timepoint

Two weeks until 1 month

### Method of measurement

Phone calls and monthly views

## Intervention groups

1

### Description

Five finger test group oral drops to 40 drops a day for  
three menstrual cycles

### Category

Treatment - Drugs

2

### Description

A placebo control group, the intervention takes place

### Category

Placebo

## Recruitment centers

1

### Recruitment center

#### Name of recruitment center

Girls' dormitory University of Shahid Chamran

#### Full name of responsible person

Parvane Mousavi

#### Street address

Girls' dormitory University Of Shahid Chamran, Esfand  
Street, Golestan

#### City

Ahvaz

## Sponsors / Funding sources

1

### Sponsor

#### Name of organization / entity

Ahvaz University Of Medical Sciences

#### Full name of responsible person

Kouroosh Zare

#### Street address

Esfand Street, Golestan

#### City

Ahvaz

### Grant name

**Grant code / Reference number**  
**Is the source of funding the same sponsor organization/entity?**

Yes

**Title of funding source**  
Ahvaz 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**  
Ahvaz University of Medical Sciences

**Full name of responsible person**  
Parvaneh Mousavi

**Position**  
Master of Science in Nursing / Academic

**Other areas of specialty/work**

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## Person responsible for scientific inquiries

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Msc Obstetrics

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## Person responsible for updating data

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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*