

Clinical Trial Protocol

Iranian Registry of Clinical Trials

07 Jul 2026

The effect of acupressure on improving quality of sleep and fatigue in female students with Premenstrual Syndrome (PMS)

Protocol summary

Summary

This is a randomized controlled trial which the primary outcomes measure include reducing fatigue and increasing quality of sleep in female students with Premenstrual Syndrome (PMS). Census sampling method will be used to screen PMS in the students of Islamic Azad University and Payam e Noor University in Bandar Genaveh, Bushehr Province. Then, 140 students with PMS and the inclusion criteria will be selected consecutively and randomly assigned to two groups of experiment (intervention) and control (with Blocking Randomization Method). The inclusion criteria include: students with fatigue and sleep quality score of 31 and 5, respectively; regular menstruation of 24-35 days; age of 18-35 years old; not married; no physical and mental diseases; no medication or traditional treatment for controlling PMS; The exclusion criteria consist no interest for participation in the trial anymore; performing incomplete acupressure for less than 6 consecutive days for any reason; irregular menses; get married during the trial; pharmacological or non-pharmacological treatments for PMS during the trial. For data collection, a self report questionnaire will be used for Premenstrual Syndrome diagnosis. Fatigue Severity Scale (FSS) and Pittsburg Sleep Quality Index (PSQI) will be used to measure the main dependent variables in the study. The independent variable will be acupressure on the points of ST36 - below the knee, in the anterior boarder of tibia and HT7 in the hands, at the wrist crease, on the radial side of the flexor carpi ulna is tendon, between the ulna and the pisiform bones. The intervention (acupressure) will be applied for 5 minutes on each points of the body. In other words, the total duration of intervention will be 20 minutes per day for 5-10 days premenstrual period in two consecutive cycles. The procedure of acupressure will be confirmed if subjects felt sore, numb, heavy, distended, and/or warm as a result of acupressure massage.

General information

Acronym

PMS

IRCT registration information

IRCT registration number: **IRCT201206126247N6**

Registration date: **2012-10-11, 1391/07/20**

Registration timing: **retrospective**

Last update:

Update count: **0**

Registration date

2012-10-11, 1391/07/20

Registrant information

Name

Farideh Bastani

Name of organization / entity

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Recruitment status

Recruitment complete

Funding source

Vice Chancellor for Research, Tehran University of Medical Sciences

Expected recruitment start date

2012-06-20, 1391/03/31

Expected recruitment end date

2012-09-20, 1391/06/30

Actual recruitment start date

empty

Actual recruitment end date

empty

Trial completion date

empty

Scientific title

The effect of acupressure on improving quality of sleep and fatigue in female students with Premenstrual Syndrome (PMS)

Public title

Acupressure, sleep and fatigue in students with Premenstrual Syndrome

Purpose

Treatment

Inclusion/Exclusion criteria

Inclusion criteria: students with fatigue and sleep quality score of 31 and 5, respectively; regular menstruation of 24-35 days; age between 18-35 years old; not married; no physical and mental diseases; no medication or traditional treatment for controlling Premenstrual Syndrome (PMS); Exclusion criteria: Unwillingness to continue participation in the trial; performing incomplete acupressure for less than 6 consecutive days for any reason; irregular menstruation; get married during the trial; pharmacological or non-pharmacological treatments for Premenstrual Syndrome during the trial.

Age

From **18 years** old to **35 years** old

Gender

Female

Phase

1

Groups that have been masked

No information

Sample size

Target sample size: **140**

Randomization (investigator's opinion)

Randomized

Randomization description

Blinding (investigator's opinion)

Not blinded

Blinding description

Placebo

Not used

Assignment

Parallel

Other design features

This is an experimental study.

Secondary Ids

empty

Ethics committees

1

Ethics committee

Name of ethics committee

The Ethics Committee of Tehran University of Medical Sciences

Street address

Keshavarz Boulevard, Ghods St.

City

Tehran

Postal code

Approval date

2012-06-20, 1391/03/31

Ethics committee reference number

91/N/130/154

Health conditions studied

1

Description of health condition studied

Premenstrual syndrome

ICD-10 code

N94.3

ICD-10 code description

Premenstrual Tension Syndrome

Primary outcomes

1

Description

Sleep Quality and Fatigue

Timepoint

pre and post intervention (before and after acupressure)

Method of measurement

Fatigue Severity Scale (FSS) and Pittsburg Sleep Quality Index (PSQI)

Secondary outcomes

empty

Intervention groups

1

Description

In this study, the intervention would be acupressure which will be applied for control of the PMS symptoms of fatigue and quality of sleep on the experiment group.

Category

Other

2

Description

The control group will be students who not exposed to acupressure or any interventions for the control of the PMS symptoms (fatigue and sleep quality).

Category

N/A

Recruitment centers

1

Recruitment center

Name of recruitment center

Islamic Azad University and Payam e Noor University of Bandar Genaveh

Full name of responsible person

Bahare Mohammad Salehi

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Bushehr Province, Bander Genevah, Saheli St, opposit to Shahr e Shadi, Islamic Azad University of Bandar Genaveh

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Full name of responsible person

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Keshavarz, Boulevard, Ghods St.

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Grant name**Grant code / Reference number**

91-02-28-17955

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

Yes

Title of funding source

Vice Chancellor for Research, Tehran 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**

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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