

Clinical Trial Protocol

Iranian Registry of Clinical Trials

11 Jun 2026

The effectiveness of consultation based on mindfulness on health promoting behaviors of adolescents with premenstrual syndrome

Protocol summary

Study aim

Determining the effect of counseling with a mindfulness approach on health-promoting behaviors in adolescents with premenstrual syndrome

Design

By cluster sampling method, 4 schools randomly selected, 2 schools of control group and 2 schools of intervention group and among the students who had the criteria to enter the research, 46 people selected by random method.

Settings and conduct

Two control group schools and two intervention group schools and then eligible students are selected with informed consent

Participants/Inclusion and exclusion criteria

Criteria for entering the study: At least 1 year after the onset of the first menstrual period, regular periods with normal intervals in each period (21-35 days), high school girls aged 17-15 years, with moderate to severe premenstrual syndrome (Score 19 or more in the PSST questionnaire, consent to participate in the intervention, as well as criteria for withdrawal from this study. Irregular menstrual cycle, use of hormonal compounds, chronic mental and physical illness, any medication to relieve the symptoms of premenstrual syndrome, Being under any kind of stressful environmental and psychotherapeutic events and taking psychiatric medications and single parent families

Intervention groups

The sample size was 92 people (46 people in each group) who were randomly selected from the criteria criteria and the intervention group saw mindfulness exercises in 8 sessions and the control group did not see any training.

Main outcome variables

Health promotion; premenstrual syndrome

General information

Reason for update

Acronym

IRCT registration information

IRCT registration number: **IRCT20200308046720N1**
Registration date: **2020-06-08, 1399/03/19**
Registration timing: **retrospective**

Last update: **2020-06-08, 1399/03/19**

Update count: **0**

Registration date

2020-06-08, 1399/03/19

Registrant information

Name

Farahnaz Moradymahmodabade

Name of organization / entity

Country

Iran (Islamic Republic of)

Phone

+98 35 3232 9473

Email address

moradyfarahnaz@yahoo.com

Recruitment status

Recruitment complete

Funding source

Expected recruitment start date

2019-09-23, 1398/07/01

Expected recruitment end date

2019-10-23, 1398/08/01

Actual recruitment start date

2019-10-07, 1398/07/15

Actual recruitment end date

2019-11-06, 1398/08/15

Trial completion date

2020-01-20, 1398/10/30

Scientific title

The effectiveness of consultation based on mindfulness on health promoting behaviors of adolescents with premenstrual syndrome

Public title

The effectiveness of consultation based on mindfulness on health promoting behaviors of adolescents with premenstrual syndrome

Purpose

Education/Guidance

Inclusion/Exclusion criteria

Inclusion criteria:

At least 1 year has passed since the first menstruation
Regular periods with normal intervals in each period (21-35 days)
High school girls aged 17-15
Moderate to severe premenstrual syndrome (score 19 or higher in the PSST questionnaire)
Satisfaction with participating in the intervention

Exclusion criteria:

Irregular menstruation
Consumption of hormonal compounds
Having a chronic mental and physical illness
Medication to relieve the symptoms of premenstrual syndrome
Being under any kind of environmental stressful event
Psychotherapy and the use of psychiatric drugs
Single parent families

Age

From **15 years** old to **17 years** old

Gender

Female

Phase

2

Groups that have been masked

- Participant

Sample size

Target sample size: **92**

Actual sample size reached: **92**

Randomization (investigator's opinion)

N/A

Randomization description

Blinding (investigator's opinion)

Single blinded

Blinding description

Random selection

Placebo

Not used

Assignment

Parallel

Other design features

Secondary Ids

empty

Ethics committees

1

Ethics committee

Name of ethics committee

Ethics Committee Yazd University of Medical Sciences

Street address

Saeedi Street

City

Meybod

Province

Yazd

Postal code

8961974360

Approval date

2020-02-20, 1398/12/01

Ethics committee reference number

IR.SSU.REC.1398.209

Health conditions studied

1

Description of health condition studied

Health promotion, premenstrual syndrome

ICD-10 code

N94.3

ICD-10 code description

Premenstrual tension syndrome

Primary outcomes

1

Description

Behaviors promoting the health of adolescents with PMS

Timepoint

Intervention before, after and one month later

Method of measurement

Walker Health Promotion Lifestyle Questionnaire

2

Description

Behaviors promoting the health of adolescents with PMS

Timepoint

Intervention before, after and one month later

Method of measurement

Walker Health Promotion Lifestyle Questionnaire

Secondary outcomes

1

Description

Premenstrual Syndrome

Timepoint

Before, after and a month after the intervention

Method of measurement

PSST questionnaire

Intervention groups

1

Description

Intervention group:46 Meybod high school girls who meet the criteria for entering the study and their pms were confirmed and randomly selected by completing two consecutive courses of the psst questionnaire one week before menstruation. The present study is a quasi-

experimental study that will be conducted in parallel with the previous and subsequent methods. In this study, the intervention group received 8 counseling sessions with a mindfulness approach on a weekly basis for 5.1-2 hours and evaluated the health-promoting behaviors of the intervention group at the beginning of the study, the end of the eighth week, and the twelfth week. In addition to evaluating health-promoting behaviors, their premenstrual syndrome is assessed with a psst questionnaire. Counseling Content: Week 1: Automated Guidance, Week 2: Dealing with Obstacles and More Control Reaction to Daily Events, Week 3: Breathing with Mind Presence, Week 4: Present Present, Week 5: Acceptance and Permission, Week 6: Thoughts are not facts, Week 7: Self-care, Week 8: Using what you have learned

Category

Behavior

2**Description**

Control group: Meybod's 46 high school girls met the criteria for entering the study, and their pms were confirmed and randomly selected by completing two consecutive courses of the psst questionnaire one week before menstruation. The present study is a quasi-experimental study that will be conducted in parallel with the previous and subsequent methods. In this research, the control group does not see any training and after the end of the research, they will receive all the training of the intervention group. Behavioral health promotion behaviors are assessed at the beginning of the study, at the end of the eighth week, and at the twelfth week. In addition to evaluating health-promoting behaviors, their premenstrual syndrome is assessed with a psst questionnaire.

Category

Behavior

Recruitment centers**1****Recruitment center****Name of recruitment center**

Meybod High Schools

Full name of responsible person

Farahnaz Moradi

Street address

Saeedi street

City

Meybod

Province

Yazd

Postal code

8961974360

Phone

+98 35 3232 9473

Email

moradyfarahnaz@yahoo.com

Sponsors / Funding sources**1****Sponsor****Name of organization / entity**

Yazd University of Medical Sciences

Full name of responsible person

Dr. Massoud Mirzaei

Street address

Commander Fallahi

City

Yazd

Province

Yazd

Postal code

8961974360

Phone

+98 35 3726 3733

Email

dvc.research@ssu.ac.ir

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

Yes

Title of funding source

Yazd 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**

Yazd University of Medical Sciences

Full name of responsible person

Farahnaz Moradymahmodabade

Position

Midwif

Latest degree

Bachelor

Other areas of specialty/work

Midwifery

Street address

Saeedi Street

City

Meybod

Province

Yazd

Postal code

8961974360

Phone

+98 35 3232 9473

Fax

Email

moradyfarahnaz@yahoo.com

Person responsible for scientific inquiries

Contact

Name of organization / entity

Yazd University of Medical Sciences

Full name of responsible person

Farahnaz Moradymahmodabade

Position

Midwif

Latest degree

Bachelor

Other areas of specialty/work

Midwifery

Street address

Saeedi Street

City

Meybod

Province

Yazd

Postal code

8961974360

Phone

+98 35 3232 9473

Fax

Email

moradyfarahnaz@yahoo.com

Person responsible for updating data

Contact

Name of organization / entity

Yazd University of Medical Sciences

Full name of responsible person

Farahnaz Moradymahmodabade

Position

Midwif

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Other areas of specialty/work

Midwifery

Street address

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Province

Yazd

Postal code

8961974360

Phone

+98 35 3232 9473

Fax

Email

moradyfarahnaz@yahoo.com

Sharing plan

Deidentified Individual Participant Data Set (IPD)

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

Study Protocol

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

Statistical Analysis Plan

No - There is not a plan to make this available

Informed Consent Form

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

Clinical Study Report

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

Analytic Code

Not applicable

Data Dictionary

Not applicable