

Clinical Trial Protocol

Iranian Registry of Clinical Trials

04 Jul 2026

Evaluation of the impact of dry cupping on primary dysmenorrhea among the students residing in Qazvin University of Medical Sciences dormitories: Random controlled clinical trial

Protocol summary

Summary

Objective: Evaluation of the impact of dry cupping on primary dysmenorrhea among the students residing in Qazvin University of Medical Sciences dormitories.
Design: Random controlled clinical trial, phase II trial.
The method of studying, the conditions of inclusion: single women in the range of 35-19 years old; living in students dormitories of Qazvin University of Medical Sciences; with regular menstrual cycles; moderate to severe primary dysmenorrhea; lack of any known chronic diseases like diabetes; the absence of symptoms like burning, itching and abnormal vaginal discharge; no hormonal problems; no history of pelvic disease, fibroids and tumors; no usage of certain medications and no stressful factors (separation of parents, death of first-degree relatives and so on) in the last six months. Major excluding conditions: being under the age of 19 years and more than 35 years; irregular menstruation; menstrual without pain; mild primary dysmenorrhea; secondary dysmenorrhea; chronic disease; pelvic disease; being under the stressors in the past six months. Sample size: two of cupping therapy groups (test) and control group (75 patients in each group). Interventions: hundred and fifty people entered the study and were randomly divided into two experimental and control groups. We use dry cupping for 3 menstrual periods at one time and then rotating within 3 days before the onset of menstruation (as prophylaxis) until end of menstruation in experimental group and no intervention in control group. During three consecutive cycle, severity of dysmenorrhea will be assessed with verbal multidimensional scoring system. The main outcome variables: the severity and duration of primary dysmenorrhea, symptoms of dysmenorrhea.

General information

Acronym

IRCT registration information

IRCT registration number: **IRCT2015120125320N1**
Registration date: **2017-02-18, 1395/11/30**
Registration timing: **retrospective**

Last update:

Update count: **0**

Registration date

2017-02-18, 1395/11/30

Registrant information

Name

Maryam Taherpour

Name of organization / entity

Qazvin University of Medical Sciences, Qazvin, Iran

Country

Iran (Islamic Republic of)

Phone

+98 28 3323 7268

Email address

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Recruitment status

Recruitment complete

Funding source

Vice chancellor for research Qazvin University of Medical Sciences

Expected recruitment start date

2016-04-03, 1395/01/15

Expected recruitment end date

2016-12-19, 1395/09/29

Actual recruitment start date

empty

Actual recruitment end date

empty

Trial completion date

empty

Scientific title

Evaluation of the impact of dry cupping on primary dysmenorrhea among the students residing in Qazvin University of Medical Sciences dormitories: Random controlled clinical trial

Public title

Evaluation of the impact of dry cupping on primary dysmenorrhea among the students residing in Qazvin University of Medical Sciences dormitories.

Purpose

Treatment

Inclusion/Exclusion criteria

Inclusion criteria: single women in the range of 35-19 years old; living in students dormitories of Qazvin University of Medical Sciences; with regular menstrual cycles; moderate to severe primary dysmenorrhea; lack of any known chronic diseases like diabetes; the absence of symptoms like burning, itching and abnormal vaginal discharge; no hormonal problems; no history of pelvic disease, fibroids and tumors; no usage of certain medications and no stressful factors (separation of parents, death of first-degree relatives and so on) in the last six months. Major excluding conditions: being under the age of 19 years and more than 35 years; irregular menstruation; menstrual without pain; mild primary dysmenorrhea; secondary dysmenorrhea; chronic disease; pelvic disease; being under the stressors in the past six months.

Age

From **19 years** old to **35 years** old

Gender

Female

Phase

2

Groups that have been masked

No information

Sample size

Target sample size: **150**

Randomization (investigator's opinion)

Randomized

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

Not blinded

Blinding description**Placebo**

Not used

Assignment

Parallel

Other design features**Secondary Ids**

empty

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

Ethics Committee of Qazvin University of Medical Sciences

Street address

Qazvin University of Medical Sciences , Shahid Bahonar Blvd , Qazvin

City

Qazvin

Postal code**Approval date**

2015-11-08, 1394/08/17

Ethics committee reference number

IR.QUMS.REC.1394.193

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

Primary dysmenorrhoea

ICD-10 code

N 94.4

ICD-10 code description

Primary dysmenorrhoea

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

severity and duration of primary dysmenorrhea

Timepoint

Before the intervention, one month, two months, three months after the intervention in case and control groups

Method of measurement

Verbal multidimensional scoring system

Secondary outcomes**1****Description**

symptoms of dysmenorrhea

Timepoint

Before the intervention, one month, two months, three months after the intervention in case and control groups

Method of measurement

Verbal multidimensional scoring system

Intervention groups**1****Description**

3 days before the onset of menstruation (as prophylaxis) until end of menstruation dry cupping for 3 menstrual periods at a time and rotations, use of the experimental group and the control group will not be any interference. In dry cupping, we are using from special cups to size 5

to 7 centimeters. Cotton soaked in alcohol that wrapped to a forceps Ignited made and in a rapid move take away inside the Cup and we're warm air inside the cup and Immediately putting up tops the lumbar sides spine and suprapubic for 10-15 minutes.

Category

Treatment - Other

2**Description**

In the control group does not take place intervention

Category

Treatment - Other

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

Qazvin University of Medical Sciences dormitories

Full name of responsible person

MaryamTaherpour

Street address

Golestan Enghelab Dormitory,Shahid Bahonar Blvd,
Qazvin

City

Qazvin

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

Vice chancellor for research, Qazvin University of
Medical Sciences

Full name of responsible person

Taghi Naserpour

Street address

Qazvin University of Medical Sciences , Shahid
Bahonar Blvd , Qazvin

City

Qazvin

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

Yes

Title of funding source

Vice chancellor for research, Qazvin 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**

Qazvin University of Medical Sciences, Qazvin, Iran

Full name of responsible person

Maryam Taherpour

Position

Master of Sciences in Midwifery/Scientific board

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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