

# Clinical Trial Protocol

## Iranian Registry of Clinical Trials

10 Jun 2026

### Effect of *Stachys lavandulifolia* on relief of primary dysmenorrheal

#### Protocol summary

##### Summary

Introduction: primary dysmenorrhea is a common symptom in women of menstrual age and it occurs between 60-93 percent of female. This symptom usually begins at the start of menstruation, continues for a few days, and is characterized by pain that radiates from the lower abdomen to the inner thighs, is usually limited to the first 48 or 72 hours of menstruation. CAM treatments for dysmenorrhea that have been studied include transcutaneous electrical nerve stimulation (TENS), acupuncture, acupressure, spinal manipulation, behavioral interventions, and herbal and dietary therapies. This study is to evaluate the therapeutic effects of *Stachys lavandulifolia* on relief of primary dysmenorrhea. Method: participants were female students of Touyserkan Azad University in western Iran. The students were randomly assigned into control (45) and experimental (45) groups. Initially, the study population was requested to consume the *Stachys lavandulifolia* flowers at a dose of 10 gr of brewed powder three time a day and for a total of 5 days (2 days before pain to 3 days after pain), based on a common traditional administration. Later, the members of both groups were followed up for three cycles.

#### General information

##### Acronym

##### IRCT registration information

IRCT registration number: **IRCT201312318360N3**

Registration date: **2014-01-04, 1392/10/14**

Registration timing: **retrospective**

Last update:

Update count: **0**

##### Registration date

2014-01-04, 1392/10/14

##### Registrant information

Name

Ensiyeh Jenabi

##### Name of organization / entity

Islamic Azad university

##### Country

Iran (Islamic Republic of)

##### Phone

+98 81 1822 1439

##### Email address

e.jenabi@tuyiau.ac.ir

##### Recruitment status

**Recruitment complete**

##### Funding source

Touyserkan Azad University

##### Expected recruitment start date

2012-08-22, 1391/06/01

##### Expected recruitment end date

2012-11-21, 1391/09/01

##### Actual recruitment start date

empty

##### Actual recruitment end date

empty

##### Trial completion date

empty

##### Scientific title

Effect of *Stachys lavandulifolia* on relief of primary dysmenorrheal

##### Public title

Effect of *Stachys lavandulifolia* on relief of primary dysmenorrheal

##### Purpose

Treatment

##### Inclusion/Exclusion criteria

Inclusion criteria: primary dysmenorrhea; being single and agree to participate in study. Exclusion criteria: history of other diseases ; irregular menstruation; drug use and mild dysmenorrhea (1-3).

##### Age

From **19 years** old to **25 years** old

##### Gender

Female

**Phase**

2-3

**Groups that have been masked**

No information

**Sample size**

Target sample size: **100**

**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**

Islamic Azad university,Touyserkan Branch

**Street address**

touyserkan-azad university

**City**

Touyserkan

**Postal code****Approval date**

2012-07-31, 1391/05/10

**Ethics committee reference number**

14568/91

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

Primary dismenorrhea

**ICD-10 code**

N94.4

**ICD-10 code description**

Primary dysmenorrhoea

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

dismenorrhea

**Timepoint**

2 day before of mens and 3 day of mens first

**Method of measurement**

Visual Analog scale

**Secondary outcomes**

empty

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

Invention group: taking Stachys lavandulifolia (10 gram flowers in 1 glass of boil water)in 2 days before of mens and 3 days mens first

**Category**

Treatment - Drugs

**2****Description**

Control group: taking placebo 2 days before mens and 3 days of mens first.

**Category**

Placebo

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

Azad university of Touyserkan

**Full name of responsible person****Street address****City**

Touyserkan

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

Touyserkan Azad university

**Full name of responsible person**

Parisa Hejrati

**Street address**

Touyserkan Azad university

**City**

Touyserkan

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

Yes

**Title of funding source**

Touyserkan Azad 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*

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## Person responsible for general inquiries

### Contact

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## Person responsible for updating data

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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*