

Clinical Trial Protocol

Iranian Registry of Clinical Trials

02 Jun 2026

The Combined Effect of black cohosh, soybean, arctium lappa L and vitex agnus- castus extracts on clinical symptoms, biomarkers of oxidative stress biomarkers and inflammation among menopausal women

Protocol summary

Study aim

Determining the combined effect of black cohosh, soybean, arctium lappa L and vitex agnus- castus extracts on clinical symptoms, biomarkers of oxidative stress biomarkers and inflammation among menopausal women

Design

Random assignment was done by the use of computer-generated random numbers. Population and sample size: In the study 54 patients who having menopause symptom among patients of eligible and referred to Kousar hospital affiliated to Qazvin University of Medical Sciences, Qazvin, Iran, will be selected.

Settings and conduct

The present study will be performed in Kousar hospital affiliated to Qazvin University of Medical Sciences, Qazvin, Iran. Patients are randomly divided into two groups receiving supplements and placebo. All people taking the supplement or placebo will take the medicine for eight weeks. Before and after the intervention, venous blood samples will be taken from the patient after 12 hours of fasting for relevant tests.

Participants/Inclusion and exclusion criteria

Inclusion criteria: fatty liver, no gastrointestinal diseases, no smoking and alcohol Exclusion criteria: thyroid disease, breast and ovary cancer , diabetes, Liver disorders, nerves, blood pressure, taking anticoagulants

Intervention groups

Patients will be assigned to receive either combined extract (intervention group: n=27) or placebo (control group: n=27).

Main outcome variables

Clinical signs including symptoms and complications of menopause respect to vasomotor, psychosocial, and physical activity, and also biomarkers of oxidative stress and inflammation in patients with menopausal symptoms will be measured before and after the intervention.

General information

Reason for update

Acronym

IRCT registration information

IRCT registration number: **IRCT20170430033730N8**

Registration date: **2021-03-11, 1399/12/21**

Registration timing: **registered_while_recruiting**

Last update: **2021-03-11, 1399/12/21**

Update count: **0**

Registration date

2021-03-11, 1399/12/21

Registrant information

Name

Seyyed Mehdi Mirhashemi

Name of organization / entity

Qazvin University of Medical Sciences

Country

Iran (Islamic Republic of)

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Recruitment status

Recruitment complete

Funding source

Expected recruitment start date

2021-02-13, 1399/11/25

Expected recruitment end date

2021-03-15, 1399/12/25

Actual recruitment start date

empty

Actual recruitment end date

empty

Trial completion date

empty

Scientific title

The Combined Effect of black cohosh, soybean, arctium lappa L and vitex agnus- castus extracts on clinical symptoms, biomarkers of oxidative stress biomarkers and inflammation among menopausal women

Public title

The Combined Effect of black cohosh, soybean, arctium lappa L and vitex agnus- castus extracts On menopause

Purpose

Treatment

Inclusion/Exclusion criteria

Inclusion criteria:

Having menopausal symptoms

Exclusion criteria:

patients who do not want to cooperate People who smoke People who drink alcohol Patients with gastrointestinal disorders Diabetic Patients Thyroid disease, breast and ovarian cancer, liver and neurological disorders, high blood pressure, taking anticoagulants

Age

From **45 years** old to **55 years** old

Gender

Female

Phase

N/A

Groups that have been masked

- Participant
- Care provider
- Investigator
- Outcome assessor

Sample size

Target sample size: **54**

Randomization (investigator's opinion)

Randomized

Randomization description

54 eligible patients are assigned to groups A and B equally using the balanced block randomization method and the online system sealedenvelope.com. The relevant list is generated using blocks with variable sizes of 6 and 8. The created list consists of eight blocks, of which five blocks are 6 and three blocks are 8. The order of the blocks and the people in each block is completely random and each person is assigned a unique code. 54 similar envelopes are selected. A unique code is registered on each package based on the mentioned list. Depending on the study groups, supplement (A) or placebo (B) is included. The envelopes are closed and each envelope is assigned to one participant.

Blinding (investigator's opinion)

Double blinded

Blinding description

Placebo is used for blinding. Supplements and placebo are poured into containers that are similar in size, shape and appearance. Instead of group names, each container is assigned a unique code. The researchers and the patient are blind until the final analysis.

Placebo

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

Shahid Bahonar Boulevard

City

Qazvin

Province

Qazvin

Postal code

34197-59811

Approval date

2020-02-29, 1398/12/10

Ethics committee reference number

IR.QUMS.REC.1398.372

Health conditions studied

1

Description of health condition studied

menopause

ICD-10 code

N95

ICD-10 code description

Menopausal and other perimenopausal disorders

Primary outcomes

1

Description

Clinical symptoms

Timepoint

Beginning and end of the study

Method of measurement

Quality of life questionnaire for menopausal women

Secondary outcomes

1

Description

Glutathione

Timepoint

Beginning and end of the study

Method of measurement

Spectrophotometry

2

Description

MDA

Timepoint

Beginning and end of the study

Method of measurement

Spectrophotometry

3

Description

TNF- α

Timepoint

Beginning and end of the study

Method of measurement

ELISA

4

Description

IL-8

Timepoint

Beginning and end of the study

Method of measurement

ELISA

5

Description

Gonadotropins

Timepoint

Beginning and end of the study

Method of measurement

ELISA

6

Description

E2

Timepoint

Beginning and end of the study

Method of measurement

ELISA

Intervention groups

1

Description

Intervention group: will receive two capsules of 550 mg of combined extract(Black cohosh, Soybean, Sweet potato, Greater burdock, Vitex) twice daily for 8 weeks. The capsules are manufactured by Barij Essence Pharmaceutical Company, Kashan, Iran.

Category

Treatment - Drugs

2

Description

Control group: Will take two capsules of 550 mg of placebo twice daily for 8 weeks. The Placebo are manufactured by Barij Essence Pharmaceutical Company, Kashan, Iran.

Category

Placebo

Recruitment centers

1

Recruitment center

Name of recruitment center

Kousar Hospital

Full name of responsible person

Seyyed Mehdi Mirhashemi

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Sponsors / Funding sources

1

Sponsor

Name of organization / entity

Qazvin University of Medical Sciences

Full name of responsible person

Mohammad-Mehdi Emamjomeh

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Grant name

Grant code / Reference number

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

Yes

Title of funding source

Qazvin 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
Qazvin University of Medical Sciences
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Professor
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Sharing plan

Deidentified Individual Participant Data Set (IPD)

No - There is not a plan to make this available

Justification/reason for indecision/not sharing IPD

There is no further information

Study Protocol

No - There is not a plan to make this available

Statistical Analysis Plan

Not applicable

Informed Consent Form

No - There is not a plan to make this available

Clinical Study Report

Not applicable

Analytic Code

Not applicable

Data Dictionary

Not applicable