

Clinical Trial Protocol

Iranian Registry of Clinical Trials

11 Sep 2026

Clinical trial of the effect of vitamin D supplementation compared with the placebo on treatment and metabolic profiles in patients with endometrial hyperplasia

Protocol summary

Study aim

The aim of this study is to determine the effects of vitamin D supplementation on treatment and metabolic profiles including insulin resistance and lipid profiles in patients with endometrial hyperplasia.

Design

Study design: Parallel double-blind (both patients and researchers) randomized controlled clinical trial. Random assignment will be done by the use of computer-generated random numbers.

Settings and conduct

Population and sample size: 60 patients with endometrial hyperplasia of eligible and referred to Naghavi Clinic affiliated to Kashan University of Medical Sciences, Kashan, Iran in the study will be selected.

Participants/Inclusion and exclusion criteria

Inclusion criteria: Patients with endometrial hyperplasia and aged 35-55 years old will be included in this study. Exclusion criteria: Menopausal women, history of cardiovascular disease, diabetes mellitus, fasting plasma glucose (FPG) > 126 mg/dL, consumption of antihyperglycemic agents including metformin, triglycerides levels > 500 mg/dL, systolic blood pressure $\geq$ 160 mmHg, diastolic blood pressure $\geq$ 110 mmHg and untreated thyroid disease

Intervention groups

Intervention: Patients will be assigned to receive either vitamin D (n=30) or placebo (n=30). Vitamin D and placebos capsules are similar in shape and size.

Main outcome variables

Outcomes: Response to treatment (primary outcomes), and insulin metabolism parameters, lipid and metabolic profiles (secondary outcomes) will be quantified at the study baseline and end-of-trial.

General information

Reason for update

Acronym

IRCT registration information

IRCT registration number: **IRCT201606275623N86**

Registration date: **2016-07-01, 1395/04/11**

Registration timing: **retrospective**

Last update: **2019-09-22, 1398/06/31**

Update count: **1**

Registration date

2016-07-01, 1395/04/11

Registrant information

Name

Zatollah Asemi

Name of organization / entity

Kashan University of Medical Sciences

Country

Iran (Islamic Republic of)

Phone

+98 36 1534 3570

Email address

asemi_z@kaums.ac.ir

Recruitment status

Recruitment complete

Funding source

Vice chancellor for research, Kashan University of Medical Sciences

Expected recruitment start date

2016-05-25, 1395/03/05

Expected recruitment end date

2016-06-24, 1395/04/04

Actual recruitment start date

empty

Actual recruitment end date

empty

Trial completion date
empty

Scientific title
Clinical trial of the effect of vitamin D supplementation compared with the placebo on treatment and metabolic profiles in patients with endometrial hyperplasia

Public title
Effect of supplementation in treatment of patients with endometrial hyperplasia

Purpose
Treatment

Inclusion/Exclusion criteria
Inclusion criteria:
 Aged 35 to 55 years Patients with endometrial hyperplasia informed consent
Exclusion criteria:
 Menopausal women History of cardiovascular disease Diabetes mellitus Fasting plasma glucose (FPG) > 126 mg/dL Consumption of antihyperglycemic agents including metformin Triglycerides levels > 500 mg/dL Systolic blood pressure ≥ 160 mmHg Diastolic blood pressure ≥ 110 mmHg Untreated thyroid disease

Age
From **35 years** old to **55 years** old

Gender
Female

Phase
3

Groups that have been masked

- Participant
- Investigator
- Outcome assessor

Sample size
Target sample size: **60**

Randomization (investigator's opinion)
Randomized

Randomization description
At study baseline and after stratification for pre-intervention BMI and age, subjects will be randomly divided into two groups to take vitamin D supplementation (n=30) or placebo (n=30). Randomization will be done by the use of computer-generated random numbers.

Blinding (investigator's opinion)
Double blinded

Blinding description
Participants, investigators or the assessors of the outcomes are unaware of the study groups.

Placebo
Used

Assignment
Parallel

Other design features

Secondary Ids
empty

Ethics committees

1

Ethics committee

Name of ethics committee

Ethics committee of Kashan University of Medical Sciences

Street address

Ghotbe Ravandi Boulevard, Kashan

City

Kashan

Province

Isfahan

Postal code

1771844351

Approval date

2016-05-24, 1395/03/04

Ethics committee reference number

IR.Kaums.REC.1395.19

Health conditions studied

1

Description of health condition studied

Endometrial hyperplasia

ICD-10 code

E11.2

ICD-10 code description

Endometrial hyperplasia

Primary outcomes

1

Description

Response to treatment

Timepoint

At the beginning of the study and after 12 weeks of intervention

Method of measurement

Sonography

Secondary outcomes

1

Description

Insulin

Timepoint

At the beginning of the study and after 12 weeks of intervention

Method of measurement

Elisa kit

2

Description

Triglycerides

Timepoint

At the beginning of the study and after 12 weeks of intervention
Method of measurement
Enzymatic kit

3

Description
Cholesterol
Timepoint
At the beginning of the study and after 12 weeks of intervention
Method of measurement
Enzymatic kit

4

Description
HDL
Timepoint
At the beginning of the study and after 12 weeks of intervention
Method of measurement
Enzymatic kit

5

Description
Total antioxidant
Timepoint
At the beginning of the study and after 12 weeks of intervention
Method of measurement
Spectrophotometry

6

Description
Glutathione
Timepoint
At the beginning of the study and after 12 weeks of intervention
Method of measurement
Spectrophotometry

7

Description
Malondialdehyde
Timepoint
At the beginning of the study and after 12 weeks of intervention
Method of measurement
Spectrophotometry

8

Description
Nitric oxide
Timepoint
At the beginning of the study and after 12 weeks of intervention
Method of measurement

Spectrophotometry

9

Description
Insulin resistance
Timepoint
At the beginning of the study and after 12 weeks of intervention
Method of measurement
Using HOMA formula

10

Description
VLDL cholesterol
Timepoint
At the beginning of the study and after 12 weeks of intervention
Method of measurement
Enzymatic kit

Intervention groups

1

Description
Intervention group: 50000 IU vitamin D (Zahravi, Tabriz, Iran), once a week, for 12 weeks orally.
Category
Treatment - Drugs

2

Description
Control group: Placebo (Barij Essence, Kashan, Iran), once a week, for 12 weeks orally.
Category
Treatment - Drugs

Recruitment centers

1

Recruitment center
Name of recruitment center
Naghavi Clinic
Full name of responsible person
Zatollah Asemi
Street address
Shahid Rajaei Avenue, Kashan
City
Kashan
Province
Isfahan
Postal code
1771844351
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+98 31 5546 3378
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asemi_z@kaums.ac.ir

Sponsors / Funding sources

1

Sponsor

Name of organization / entity

Vice chancellor for research, Kashan University of Medical Sciences

Full name of responsible person

Gholamali Hamidi

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Ghotbe Ravandi Boulevard, Kashan

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Gholamali_h@kaums.ac.ir

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, Kashan 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**

Kashan University of Medical Sciences

Full name of responsible person

Zatollah Asemi

Position

PhD of Nutrition

Latest degree

Ph.D.

Other areas of specialty/work

Nutrition

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Web page address

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

Undecided - It is not yet known if there will be 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

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

Data Dictionary

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