

# Clinical Trial Protocol

## Iranian Registry of Clinical Trials

11 Jun 2026

### Clinical trial of the effect of Q10 supplementation compared with the placebo on hormonal profiles, inflammatory factors and oxidative stress biomarkers in women with polycystic ovary syndrome

#### Protocol summary

##### Study aim

Objective: The aim of this study is to determine the effects of Q10 supplementation on hormonal profiles, inflammatory factors and oxidative stress biomarkers in patients with polycystic ovary syndrome

##### Design

Study design: Randomized double-blind placebo-controlled trial. Randomization will be done by the use of computer-generated random numbers. Patients will be assigned into two groups to receive Q10 supplement (n=30) or placebo (n=30).

##### Settings and conduct

Among patients with polycystic ovary syndrome referred to Akbarabadi Clinic affiliated to Iran University of Medical Sciences, 60 patients will be selected according to inclusion and exclusion criteria. Participants, investigators or the assessors of the outcomes are unaware of the study groups. Supplements and placebos are similar in shape and size. Fasting blood samples will be taken at baseline and 12 weeks after the intervention. At the beginning and the end of the intervention: 12 weeks.

##### Participants/Inclusion and exclusion criteria

Inclusion criteria: Patients with polycystic ovary syndrome aged 18 to 40 years. Exclusion criteria: Unwillingness to cooperate.

##### Intervention groups

Intervention group: 100 mg Q10 (Nature, New York, USA), once a day, for 12 weeks orally. Control group: Placebo (Barij Essence, Kashan, Iran), once a day, for 12 weeks orally.

##### Main outcome variables

Outcomes: Total testosterone and hs-CRP (primary outcome) and biomarkers of oxidative stress and mental health parameters (secondary outcomes) will be quantified at study baseline and end-of-trial.

#### General information

##### Reason for update

##### Acronym

##### IRCT registration information

IRCT registration number: **IRCT20170513033941N36**

Registration date: **2018-06-25, 1397/04/04**

Registration timing: **registered\_while\_recruiting**

Last update: **2018-06-25, 1397/04/04**

Update count: **0**

##### Registration date

2018-06-25, 1397/04/04

##### Registrant information

##### Name

Mohammadreza Sharif

##### Name of organization / entity

##### Country

Iran (Islamic Republic of)

##### Phone

+98 31 5546 3378

##### Email address

ostadmohammadi-vr@kaums.ac.ir

##### Recruitment status

**Recruitment complete**

##### Funding source

##### Expected recruitment start date

2018-06-14, 1397/03/24

##### Expected recruitment end date

2018-07-01, 1397/04/10

##### Actual recruitment start date

empty

##### Actual recruitment end date

empty

##### Trial completion date

empty

## Scientific title

Clinical trial of the effect of Q10 supplementation compared with the placebo on hormonal profiles, inflammatory factors and oxidative stress biomarkers in women with polycystic ovary syndrome

## Public title

Effect of Q10 supplementation in treatment of women with polycystic ovary syndrome

## Purpose

Treatment

## Inclusion/Exclusion criteria

### Inclusion criteria:

Inclusion criteria: Patients with polycystic ovary syndrome. Individuals aged 18 to 40 years.

### Exclusion criteria:

Exclusion criteria: Unwillingness to cooperate.

## Age

From **18 years** old to **40 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 receive supplement or placebo. 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 Iran University of Medical Sciences

##### Street address

Iran University of Medical Sciences, Hemmat Highway, Tehran

##### City

Tehran

##### Province

Tehran

##### Postal code

1449614535

##### Approval date

2018-06-13, 1397/03/23

##### Ethics committee reference number

IR.IUMS.FMD.REC.1396.9411290014

## Health conditions studied

### 1

#### Description of health condition studied

Polycystic ovary syndrome

#### ICD-10 code

E28.2

#### ICD-10 code description

Polycystic ovarian syndrome

## Primary outcomes

### 1

#### Description

Total testosterone

#### Timepoint

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

#### Method of measurement

Elisa kit

### 2

#### Description

Hs-CRP

#### Timepoint

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

#### Method of measurement

Elisa kit

## Secondary outcomes

### 1

#### Description

SHBG

#### Timepoint

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

#### Method of measurement

Elisa kit

### 2

#### Description

Malondialdehyde

### **Timepoint**

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

### **Method of measurement**

Spectrophotometry

### **3**

### **Description**

Glutathione

### **Timepoint**

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

### **Method of measurement**

Spectrophotometry

### **4**

### **Description**

Total antioxidant capacity

### **Timepoint**

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

### **Method of measurement**

Spectrophotometry

### **5**

### **Description**

Beck Depression Inventory

### **Timepoint**

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

### **Method of measurement**

Questionnaire

### **6**

### **Description**

General Health Questionnaire

### **Timepoint**

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

### **Method of measurement**

Questionnaire

## **Intervention groups**

### **1**

### **Description**

Intervention group: 100 mg Q10 (Nature, New York, USA), once a day, for 12 weeks orally.

### **Category**

Treatment - Drugs

### **2**

### **Description**

Control group: Placebo (Barij Essence, Kashan, Iran), once a day, for 12 weeks orally

### **Category**

Placebo

## **Recruitment centers**

### **1**

### **Recruitment center**

#### **Name of recruitment center**

Akbarabadi Clinic

#### **Full name of responsible person**

Dr. Maryam Karamali

#### **Street address**

Akbarabadi Hospital, Mowlavi Street, Tehran

#### **City**

Tehran

#### **Province**

Tehran

#### **Postal code**

1168743514

#### **Phone**

+98 21 5560 6034

#### **Email**

karamali.maryam2@gmail.com

## **Sponsors / Funding sources**

### **1**

### **Sponsor**

#### **Name of organization / entity**

Iran University of Medical Sciences

#### **Full name of responsible person**

Dr. Seyed Kazem Malakouti

#### **Street address**

Vice chancellor for research, Iran University of Medical Sciences, Hemmat Highway, Tehran

#### **City**

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#### **Province**

Tehran

#### **Postal code**

1168743514

#### **Phone**

+98 21 6650 9024

#### **Email**

kmalakouti@iums.ac.ir

### **Grant name**

### **Grant code / Reference number**

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

Yes

### **Title of funding source**

Iran 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

**Street address**

Ghotbe Ravandi Boulevard, Kashan

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asemi\_z@kaums.ac.ir

**Web page address**

## Person responsible for updating data

### Contact

**Name of organization / entity**

Kashan University of Medical Sciences

**Full name of responsible person**

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## Person responsible for scientific inquiries

### Contact

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Kashan University of Medical Sciences

**Full name of responsible person**

Zatollah Asemi

**Position**

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**Latest degree**

Ph.D.

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## 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