

# Clinical Trial Protocol

## Iranian Registry of Clinical Trials

07 Jun 2026

### The effect of oral sachet containing extracts of *Garcinia cambogia* and *Plantago psyllium* with capsules containing extract of Green tea and Green coffee on weight of obese and over weight people

#### Protocol summary

##### Study aim

The effect of oral sachet containing extracts of *Garcinia cambogia* and *Plantago psyllium* with capsules containing extract of Green tea and Green coffee and placebo on weight of over weight and obese people in Beheshti nutrition therapy center The effect of oral sachet containing extracts of *Garcinia cambogia* and *Plantago psyllium* with capsules containing extract of Green tea and Green coffee and placebo on blood lipid level(Cholesterol, Triglyceride and High and low density lipoprotein), fasting blood glucose, liver enzymes and high sensitive-Critical Reactive Protein and plasma total antioxidant

##### Design

Randomized clinical trial, controlled with placebo, parallel, double blind, single central

##### Settings and conduct

This study is a clinical trial on 82 patients BMI>25, aged 18 to 50 who refers to Kashan Beheshti nutrition therapy center. Individuals based on table of random numbers will be divided to one of the groups received herbal or placebo treatment plus standard treatment for 8 weeks randomly . Herbal drug and placebo are presented in the same package and the physician , patients, researcher and statistical analyst are blind .After analyzing data, codes will be decoded.

##### Participants/Inclusion and exclusion criteria

Inclusion criteria:18-50 years old.BMI>25 exclusion criteria: metabolic disorders; nutritional supplements; pregnancy; lactation; thyroid and parathyroid disease.

##### Intervention groups

Intervention group: Weight loss diet+regular physical activity+ oral sachet containing extracts of *Garcinia cambogia* and *Plantago psyllium* half an hour before meals and oral capsules containing extract of Green tea and Green coffee after each meal twice a day.each time1. for 8 weeks Control group: Weight loss

diet+regular physical activity+ oral placebo sachet half an hour before meals and placebo capsules after each meal twice a day.each time1. for 8 weeks

##### Main outcome variables

At the end of study: changes of weight: blood Cholesterol:Triglyceride: high and low density lipoprotein: fasting blood glucose: liver enzymes: high sensitive-Critical Reactive Protein and plasma total antioxidant

#### General information

##### Reason for update

##### Acronym

##### IRCT registration information

IRCT registration number: **IRCT20130211012438N25**

Registration date: **2018-02-26, 1396/12/07**

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

Last update: **2018-02-26, 1396/12/07**

Update count: **0**

##### Registration date

2018-02-26, 1396/12/07

##### Registrant information

##### Name

Mohsen Taghizadeh

##### Name of organization / entity

Kashan University of Medical Sciences

##### Country

Iran (Islamic Republic of)

##### Phone

+98 36 1555 0021

##### Email address

taghizadeh\_m@kaums.ac.ir

##### Recruitment status

**Recruitment complete**

##### Funding source

**Expected recruitment start date**

2018-02-20, 1396/12/01

**Expected recruitment end date**

2018-08-23, 1397/06/01

**Actual recruitment start date**

empty

**Actual recruitment end date**

empty

**Trial completion date**

empty

**Scientific title**

The effect of oral sachet containing extracts of Garcinia cambogia and Plantago psyllium with capsules containing extract of Green tea and Green coffee on weight of obese and over weight people

**Public title**

The effect of oral sachet containing extracts of Garcinia cambogia and Plantago psyllium with capsules containing extract of Green tea and Green coffee on obesity

**Purpose**

Treatment

**Inclusion/Exclusion criteria****Inclusion criteria:**

18-50 years old BMI>25

**Exclusion criteria:**

metabolic disorders nutritional supplements pregnancy and lactation thyroid and parathyroid disease

**Age**

From **18 years** old to **50 years** old

**Gender**

Both

**Phase**

2-3

**Groups that have been masked**

- Participant
- Care provider
- Investigator
- Outcome assessor
- Data analyser

**Sample size**

Target sample size: **82**

**Randomization (investigator's opinion)**

Randomized

**Randomization description**

Randomized based on table of random numbers

**Blinding (investigator's opinion)**

Double blinded

**Blinding description**

Coding and similar packaging. The physician, patient, researcher and statistical analyzer are blind. At the end of the study and analysis of data, codes are decoded.

**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**

Kashan University Of Medical Sciences, Kashan

**City**

Kashan

**Province**

Isfahan

**Postal code**

81151-87159

**Approval date**

2018-01-08, 1396/10/18

**Ethics committee reference number**

IR.KAUMS.NUHEPM.REC.1396.28

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

Overweight and obesity

**ICD-10 code**

E66

**ICD-10 code description**

Overweight and obesity

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

Weight changes in people with BMI greater than 25.

**Timepoint**

At the beginning of the study, the end of the fourth week, the end of the study

**Method of measurement**

digital scale

**2****Description**

lipid profile (Cholesterol, Triglyceride and High and low density lipoprotein)

**Timepoint**

Baseline and End-of-trial

**Method of measurement**

fasting blood sample

**3****Description**

fasting blood glucose

## Timepoint

Baseline and End-of-trial

## Method of measurement

fasting blood sample

## 4

### Description

liver enzymes

### Timepoint

Baseline and End-of-trial

### Method of measurement

blood sample

## 5

### Description

high sensitive-Critical Reactive Protein

### Timepoint

Baseline and End-of-trial

### Method of measurement

blood sample

## 6

### Description

plasma total antioxidant

### Timepoint

Baseline and End-of-trial

### Method of measurement

blood sample

## Secondary outcomes

empty

## Intervention groups

### 1

#### Description

Intervention group: Weight loss diet+regular physical activity+ oral sachet containing extracts of Garcinia cambogia and Plantago psyllium half an hour before meals and oral capsules containing extract of Green tea and Green coffee after each meal twice a day.each time1. for 8 weeks

#### Category

Treatment - Drugs

### 2

#### Description

Control group: Weight loss diet+regular physical activity+ oral placebo sachet half an hour before meals and placebo capsules after each meal twice a day.each time1. for 8 weeks

#### Category

Treatment - Drugs

## Recruitment centers

### 1

#### Recruitment center

##### Name of recruitment center

Kashan Beheshti nutrition therapy center

##### Full name of responsible person

Dr Mohsen Taghizadeh

##### Street address

Shahid beheshti hospital, Ghotbe Ravandi Boulevard, Kashan

##### City

Kashan

##### Province

Isfahan

##### Postal code

87159/81151

##### Phone

+98 31 5554 0026

##### Fax

+98 31 5554 8900

##### Email

mohsenta44@yahoo.com

##### Web page address

<http://www.beheshti.kaums.ac.ir/>

## Sponsors / Funding sources

### 1

#### Sponsor

##### Name of organization / entity

Kashan University of Medical Sciences

##### Full name of responsible person

Dr Gholam Ali Hamidi

##### Street address

Kashan university of Medical Science, Kashan

##### City

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##### Province

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##### Postal code

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##### Phone

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##### Email

hamidi@yahoo.com

##### Web page address

<http://kaums.ac.ir>

#### Grant name

#### Grant code / Reference number

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

Yes

#### Title of funding source

Kashan University of Medical Sciences

#### Proportion provided by this source

50

#### 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

## 2

### **Sponsor**

**Name of organization / entity**  
Barij Research Center of Medicinal Herbs

**Full name of responsible person**  
Laleh Hejazi

**Street address**  
Kashan Mashhad Ardehal Road, KM44

**City**  
Kashan, Mashhad ardehal

**Province**  
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1178

**Phone**  
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**Fax**  
+98 86 4436 2187

**Email**  
info@barijessence.com

**Web page address**  
<http://www.barijessence.com>

### **Grant name**

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

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

Yes

### **Title of funding source**

Barij Research Center of Medicinal Herbs

### **Proportion provided by this source**

50

### **Public or private sector**

Private

### **Domestic or foreign origin**

Domestic

### **Category of foreign source of funding** *empty*

### **Country of origin**

### **Type of organization providing the funding** Industry

## **Person responsible for general inquiries**

### **Contact**

**Name of organization / entity**  
Kashan University of Medical Sciences

**Full name of responsible person**  
Dr Mohsen Taghizadeh

**Position**  
Associate professor

**Latest degree**  
Ph.D.

**Other areas of specialty/work**  
Nutrition

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## **Person responsible for scientific inquiries**

### **Contact**

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**Full name of responsible person**  
Mohsen Taghizadeh

**Position**  
Associate professor

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**Other areas of specialty/work**  
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taghizadeh\_m@kaums.ac.ir

## **Person responsible for updating data**

### **Contact**

**Name of organization / entity**  
Barij Research Center of Medicinal Herbs

**Full name of responsible person**  
Mahnaz Mahlouji

**Position**  
کارشناس تحقیقات بالینی

**Latest degree**  
Bachelor

**Other areas of specialty/work**  
Midwifery

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Kashan Mashhad Ardehal Road, KM44

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**Province**  
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**Postal code**  
1178

**Phone**

+98 86 4854 0000

**Email**

mahnaz.mahlouji@yahoo.com

**Web page address**

<http://www.barijessence.com>

**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