

Clinical Trial Protocol

Iranian Registry of Clinical Trials

12 Jun 2026

Effect of oral soft gel containing lemon and cumin essential oil in various doses on weight in subjects with overweight and obesity

Protocol summary

Study aim

The aim of this study is to determine the effect of oral soft gel containing lemon and cumin essential oil in various doses on weight in subjects with BMI \geq 25.

Design

Parallel double-blind randomized controlled clinical trial.

Settings and conduct

72 subjects with overweight and obese eligible and referred to Naghavi Diet Therapy Clinic affiliated to Kashan University of Medical Sciences, Kashan, Iran in the study will be selected.

Participants/Inclusion and exclusion criteria

Inclusion criteria: Subjects aged 18 - 50 years with overweight or obesity (BMI \geq 25 kg/m²). The exclusion criteria will as follows: BMI < 25 kg/m², subjects with hypertension, thyroid, diabetes or cardiovascular disorders, pregnancy and lactation and the use of hormonal or anti-obesity medications.

Intervention groups

Subjects will be assigned to receive either oral 25 mg soft gel containing lemon and cumin essential oil (intervention group: n=24), oral 75 mg soft gel containing lemon and cumin essential oil (intervention group: n=24) or placebo (control group: n=24).

Main outcome variables

Weight, fasting plasma glucose (FPG), serum insulin, insulin resistance, oxidative stress, lipid profiles and thyroid hormones

General information

Reason for update

Due to an error, the request for an update in our website has conducted after paper published. However, the revisions were accordance with either the original approved proposal or with coordination with Vice Chancellor of Research at the University.

Acronym

IRCT registration information

IRCT registration number: **IRCT2015030812438N10**

Registration date: **2015-03-25, 1394/01/05**

Registration timing: **registered_while_recruiting**

Last update: **2020-12-22, 1399/10/02**

Update count: **1**

Registration date

2015-03-25, 1394/01/05

Registrant information

Name

Mohsen Taghizadeh

Name of organization / entity

Kashan University of Medical Sciences

Country

Iran (Islamic Republic of)

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Recruitment status

Recruitment complete

Funding source

Barij Essence Company

Expected recruitment start date

2015-03-21, 1394/01/01

Expected recruitment end date

2015-03-30, 1394/01/10

Actual recruitment start date

empty

Actual recruitment end date

empty

Trial completion date

empty

Scientific title

Effect of oral soft gel containing lemon and cumin essential oil in various doses on weight in subjects with overweight and obesity

Public title

Effect of cumin and lemon on obesity

Purpose

Treatment

Inclusion/Exclusion criteria**Inclusion criteria:**

Overweight or obesity (BMI \geq 25 kg/m²) Aged 18-50 years

Exclusion criteria:

BMI < 25 kg/m² Hypertension Thyroid disorders Diabetes Cardiovascular disorders Pregnancy and lactation The use of hormonal or anti-obesity medications

Age

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

Gender

Both

Phase

3

Groups that have been masked

- Participant
- Investigator
- Outcome assessor

Sample size

Target sample size: **72**

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 three groups to take either high-dose C. cyminum L. and lime supplements (n =24) or low-dose C. cyminum L. and lime supplements (n = 24) or placebo (n = 24). Randomization will be done by the use of computer-generated random numbers.

Blinding (investigator's opinion)

Double blinded

Blinding description

Randomization and allocation will be concealed from the researchers and participants until the final analyses are completed. Another person who is not involved in the trial and not aware of random sequences, will be assigned the participants to the numbered bottles of capsules.

Placebo

Used

Assignment

Parallel

Other design features**Secondary Ids**

empty

Ethics committees**1****Ethics committee****Name of ethics committee**

Kashan University of Medical Sciences

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Kashan

City

Bolvare Ghotbe Ravandi, Kashan

Province

Isfahan

Postal code

8715988141

Approval date

2015-02-28, 1393/12/09

Ethics committee reference number

93210

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

Overweight and Obesity

ICD-10 code

E66.9

ICD-10 code description

Obesity, unspecified

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

Weight

Timepoint

Baseline and End-of-trial

Method of measurement

scale

2**Description**

Body Mass Index

Timepoint

Baseline and End-of-trial

Method of measurement

Calculation using formula

Secondary outcomes**1****Description**

Total cholesterol

Timepoint

Baseline and End-of-trial

Method of measurement

Enzymatic kit

2**Description**

Insulin

Timepoint

Baseline and End-of-trial

Method of measurement

ELISA kit

3

Description

Triglycerides

Timepoint

Baseline and End-of-trial

Method of measurement

Enzymatic kit

4

Description

HDL-cholesterol

Timepoint

Baseline and End-of-trial

Method of measurement

Enzymatic kit

5

Description

LDL-cholesterol

Timepoint

Baseline and End-of-trial

Method of measurement

Enzymatic kit

6

Description

Glutathione

Timepoint

Baseline and End-of-trial

Method of measurement

Spectrophotometry

7

Description

Total antioxidant capacity

Timepoint

Baseline and End-of-trial

Method of measurement

Spectrophotometry

8

Description

Fasting blood sugar

Timepoint

Baseline and End-of-trial

Method of measurement

Enzymatic kit

9

Description

Insulin resistance

Timepoint

Baseline and End-of-trial

Method of measurement

Calculation using HOMA formula

Intervention groups

1

Description

Intervention group: Oral soft gel containing lemon and cumin essential oil, 25 mg, twice a day, for 8 weeks orally.

Category

Treatment - Drugs

2

Description

Intervention group: Oral soft gel containing lemon and cumin essential oil, 75 mg, twice times a day, for 8 weeks orally

Category

Treatment - Drugs

3

Description

Control group: Placebo pearl, twice times a day, for 8 weeks orally.

Category

Placebo

Recruitment centers

1

Recruitment center**Name of recruitment center**

Naghavi Clinic

Full name of responsible person

Zatollah Asemi

Street address

Shahid Rajaei Avenue, Kashan

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Sponsors / Funding sources

1

Sponsor**Name of organization / entity**

Kashan University of Medical Sciences

Full name of responsible person

Gholamali Hamidi

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Grant name
Grant code / Reference number
Is the source of funding the same sponsor organization/entity?
No
Title of funding source
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
Dr.M.Taghizadeh
Position
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Latest degree
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Person responsible for updating data

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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