

Clinical Trial Protocol

Iranian Registry of Clinical Trials

26 May 2026

The Effect of 10 weeks of high intensity interval training and green tea supplementation on serum levels of Sirtuin 1, PGC-1 α and Catalase in overweight women

Protocol summary

Summary

The aim of the present study is determining the effect of 10 weeks of high intensity interval training with consuming green tea supplementation on Sirtuin-1 (SIRT1) and Peroxisome Proliferate-Activated Receptor Gamma Co activator 1-Alpha (PGC-1 α) in overweight women. Thirty non-athlete females having overweight randomly divided into 3 consistent and equal groups including high intensity interval training along with supplements group, High intensity interval training plus placebo and supplementary group. The training program included 3 sessions in a week for 10 weeks with the intensity of 85-95% maximum heart rate. The groups consuming green tea consumed 3 tablets of 500 mg after each main meal for 10 weeks. Venous blood samples before test and post test was used for measuring SIRT1 and PGC-1 α . Body mass index, fat percentage and the weight of the participants were assessed before and after intervention.

General information

Acronym

IRCT registration information

IRCT registration number: **IRCT2015121425524N1**

Registration date: **2016-04-01, 1395/01/13**

Registration timing: **prospective**

Last update:

Update count: **0**

Registration date

2016-04-01, 1395/01/13

Registrant information

Name

Elham Ghasemi

Name of organization / entity

Birjand University

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Recruitment status

Recruitment complete

Funding source

Thesis, Vice chancellor Of Research, Birjand University

Expected recruitment start date

2016-05-21, 1395/03/01

Expected recruitment end date

2016-07-22, 1395/05/01

Actual recruitment start date

empty

Actual recruitment end date

empty

Trial completion date

empty

Scientific title

The Effect of 10 weeks of high intensity interval training and green tea supplementation on serum levels of Sirtuin 1, PGC-1 α and Catalase in overweight women

Public title

The Effect of 10 weeks of high intensity interval training and green tea supplementation on serum levels of Sirtuin 1, PGC-1 α and Catalase in overweight women

Purpose

Prevention

Inclusion/Exclusion criteria

Inclusion criteria: Lack of participation in regular physical activity; lack of any recent chronic disease; non-smoking; lack of cardiovascular disease, metabolic or respiratory diseases; not taking any herbal supplement or drug or

chemical antioxidant and extreme sports in the last 6 months. Exclusion criteria: lack of regular participation in exercise; lack of regular use of supplements; Taking herbal supplements and antioxidant drug or chemical or strenuous exercise during the study period.

Age

From **19 years** old to **30 years** old

Gender

Female

Phase

N/A

Groups that have been masked

No information

Sample size

Target sample size: **30**

Randomization (investigator's opinion)

Randomized

Randomization description

Blinding (investigator's opinion)

Single blinded

Blinding description

Placebo

Used

Assignment

Parallel

Other design features

Secondary Ids

empty

Ethics committees

1

Ethics committee

Name of ethics committee

Ethics Committee of Birjand University of Medical Sciences

Street address

Birjand University of Medical Sciences, Birjand, South Khorasan

City

Birjand

Postal code

Approval date

2016-02-08, 1394/11/19

Ethics committee reference number

ir.bums.1394.312

Health conditions studied

1

Description of health condition studied

Overweight and Obesity

ICD-10 code

E65,E66,E6

ICD-10 code description

Localized adiposity, Obesity, Other hyperalimentation, Sequelae of hyperalimentation

Primary outcomes

1

Description

Sirtuin 1

Timepoint

Baseline and after 10 weeks

Method of measurement

ELISA

2

Description

PGC-1a

Timepoint

Baseline and after 10 weeks

Method of measurement

ELISA

Secondary outcomes

1

Description

Weight

Timepoint

Baseline and after 10 weeks

Method of measurement

Digital Balance

2

Description

Body mass index

Timepoint

Baseline and after 10 weeks

Method of measurement

Formula

3

Description

Fat percentage

Timepoint

Baseline and after 10 weeks

Method of measurement

Caliper and Jackson Pollock formula

Intervention groups

1

Description

The intervention group practice with supplements of green tea: consumption 10 weeks 500 mg Dinah green tea tablets three times a day , along with intensive interval training with intensity of 85-95% of maximum heart rate

Category

Placebo

2

Description

Intervention group supplements of green tea: consumption 500 mg Dinah green tea tablets for 10 weeks in three times a day.

Category

Placebo

3

Description

The intervention group placebo: 10 weeks of taking 500 mg starch powder tablets similar to green tea tablets three times daily.

Category

Placebo

Recruitment centers

1

Recruitment center

Name of recruitment center

Birjand University

Full name of responsible person

Doctor Mohammad Ismail Afzalpour

Street address

Birjand University, Birjand, South Khorasan

City

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Sponsors / Funding sources

1

Sponsor

Name of organization / entity

Vice Chancellor Of Research, Birjand University

Full name of responsible person

Doctor Mohammad Ali Behdani

Street address

Birjand University, Birjand

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Birjand

Grant name

Grant code / Reference number

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

Yes

Title of funding source

Vice Chancellor Of Research, Birjand University

Proportion provided by this source

100

Public or private sector

empty

Domestic or foreign origin

empty

Category of foreign source of funding

empty

Country of origin

Type of organization providing the funding

empty

Person responsible for general inquiries

Contact

Name of organization / entity

Birjand University

Full name of responsible person

Elham Ghasemi

Position

PhD Student In Exercise Physiology

Other areas of specialty/work

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Web page address**Sharing plan****Deidentified Individual Participant Data Set (IPD)**

empty

Study Protocol

empty

Statistical Analysis Plan

empty

Informed Consent Form

empty

Clinical Study Report

empty

Analytic Code

empty

Data Dictionary

empty