

# Clinical Trial Protocol

## Iranian Registry of Clinical Trials

10 Jul 2026

### Changes in cardiovascular risk factors of overweight men after eight weeks supplementation of green tea and weight training

#### Protocol summary

##### Summary

The aim of the present study was to investigate the effect of eight weeks of green tea supplementation and weight training on cardiovascular risk factors in overweight men. Methods: 30 overweight male with a body mass index between 30-25 (kg/m<sup>2</sup>), were randomly divided to three groups of 10 subjects: 1. Weight training and supplementation with green tea (supplement), 2. Weight training (exercise) and 3. control group. Weight training was consisting of three sessions per week with an intensity of 60% - 80% of one repetition maximum for 8 weeks. Green tea was also used to supplement the amount of 6 grams per day, in three equal portions. Inclusion criteria included: No history of chronic disease; orthopedic problems; smoking; and taking antioxidant supplements. Exclusion criteria included: Absence of more than three sessions during practice; gastrointestinal distress and physical injuries. Body compositions of all subjects were assessed at pre-test and post-test using calipers. Venous blood samples, were used in the pre-test and post-test for analysis of total cholesterol, HDL, LDL, triglycerides and fibrinogen.

#### General information

##### Acronym

##### IRCT registration information

IRCT registration number: **IRCT2013080414267N1**

Registration date: **2014-01-20, 1392/10/30**

Registration timing: **retrospective**

Last update:

Update count: **0**

##### Registration date

2014-01-20, 1392/10/30

##### Registrant information

**Name**

Worya Tahmasebi

##### Name of organization / entity

Razi university

##### Country

Iran (Islamic Republic of)

##### Phone

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##### Email address

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##### Recruitment status

**Recruitment complete**

##### Funding source

Investigator

##### Expected recruitment start date

2013-04-09, 1392/01/20

##### Expected recruitment end date

2013-06-10, 1392/03/20

##### Actual recruitment start date

empty

##### Actual recruitment end date

empty

##### Trial completion date

empty

##### Scientific title

Changes in cardiovascular risk factors of overweight men after eight weeks supplementation of green tea and weight training

##### Public title

Effect of eight weeks of green tea supplementation on cardiovascular risk factors in overweight individuals

##### Purpose

Prevention

##### Inclusion/Exclusion criteria

Inclusion criteria: Overweight men with body mass index between 25-30 kg/m<sup>2</sup>; age range 18-27 years; no history of chronic disease; orthopedic problems; smoking and taking antioxidant supplements. Exclusion criteria: More than three absences during a training session;

gastrointestinal discomfort and physical injury.

#### Age

From **18 years** old to **27 years** old

#### Gender

Male

#### 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)

Not blinded

#### Blinding description

#### Placebo

Not used

#### Assignment

Parallel

#### Other design features

30 volunteers were randomly divided (lucky draw) to three groups of 10 subjects (10 subjects in each group).

## Secondary Ids

empty

## Ethics committees

### 1

#### Ethics committee

##### Name of ethics committee

Ethics committee of University of Kurdistan

##### Street address

University of Kurdistan, Pasdaran Street.

##### City

Sanandaj

##### Postal code

#### Approval date

2012-12-27, 1391/10/07

#### Ethics committee reference number

3/2684

## Health conditions studied

### 1

#### Description of health condition studied

Cardiovascular disease

#### ICD-10 code

I25.0

#### ICD-10 code description

Atherosclerotic cardiovascular disease

## Primary outcomes

### 1

#### Description

Cholesterol, TG, HDL and LDL

#### Timepoint

Before and after intervention

#### Method of measurement

Cholesterol, triglycerides, HDL and LDL were measured with spectrophotometer method using kits from Pars Azmun company Iran.

### 2

#### Description

Fibrinogen

#### Timepoint

Before and after intervention

#### Method of measurement

Fibrinogen concentration was measured with thrombin clotting time method using sigma kit from Kyowl Medex company, Japan.

## Secondary outcomes

empty

## Intervention groups

### 1

#### Description

Intervention group 1: Weight training and supplementation with green tea: 10 subjects did weight training for 8 weeks, 3 sessions per week, with an intensity of 60-80% of one repetition maximum. In addition, subjects who consumed a daily 6 grams of dried leaf powder, green tea (2 g powder of dried green tea leaves in 200 ml water at 80°C) for one hour before meals at 3 times (morning, afternoon and evening). Green tea was in bag (2 g) and manufactured in the Lahigan Tea Company, Iran.

#### Category

Prevention

### 2

#### Description

Intervention groups 2: Weight training, 10 participants did resistance training for 8 weeks and 3 sessions per week, with an intensity of 60-80% of one repetition maximum.

#### Category

Prevention

### 3

#### Description

Control group: 10 subjects did no exercise and supplementation and only had to do routine activities.

#### Category

Prevention

## Recruitment centers

1

### Recruitment center

**Name of recruitment center**

University of Kurdistan

**Full name of responsible person**

Mohammadzadeh Farid, MS, Instructor

**Street address**

University of Kurdistan, Pasdaran St., Sanandaj,  
Kurdistan state

**City**

Sanandaj

## Sponsors / Funding sources

1

### Sponsor

**Name of organization / entity**

Vice Chancellor for research of University of Kurdistan

**Full name of responsible person**

Abdollah Salimi

**Street address**

University of Kurdistan, Pasdaran Street

**City**

Sanandaj

**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 of University of Kurdistan

**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**

University of Kurdistan

**Full name of responsible person**

Farhad Ahmadi Kani Golzar

**Position**

MA, Lecturer

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## Person responsible for scientific inquiries

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## Person responsible for updating data

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## 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*