

Clinical Trial Protocol

Iranian Registry of Clinical Trials

22 Jul 2026

Study of Portulaca Oleracea and Cumin effects on serum paraoxanase 1 activity

Protocol summary

Summary

In this clinical trial study patients are recruited from Clinic of Internal Medicine. Patients with low density lipoprotein cholesterol (LDL-C) over than 120 mg/dl are included in the study and divided in three groups (Cumin, Purslane and lovastatin). These patients are matched for age, gender, exercise and smoking. A fasting venous blood sample is obtained from all participants before initiating treatment and a second sample after a mean of 45 days of treatment. Total cholesterol (TC), triglycerides (TG), high density lipoprotein cholesterol (HDL-C), LDL-C, apolipoprotein A1 (Apo A1), apolipoprotein B (Apo B), arylesterase activity and PON1 activity toward paraoxon and the oxidised LDL (oxLDL) are measured before and 45 days after the treatment.

General information

Acronym

IRCT registration information

IRCT registration number: **IRCT138902063806N1**

Registration date: **2010-04-21, 1389/02/01**

Registration timing: **retrospective**

Last update:

Update count: **0**

Registration date

2010-04-21, 1389/02/01

Registrant information

Name

Effat Farrokhi

Name of organization / entity

Shahrekord University of Medical Sciences

Country

Iran (Islamic Republic of)

Phone

+98 38 1334 6692

Email address

farrokhi@skums.ac.ir

Recruitment status

Recruitment complete

Funding source

Shahrekord university of medical sciences

Expected recruitment start date

2010-04-21, 1389/02/01

Expected recruitment end date

2010-06-22, 1389/04/01

Actual recruitment start date

empty

Actual recruitment end date

empty

Trial completion date

empty

Scientific title

Study of Portulaca Oleracea and Cumin effects on serum paraoxanase 1 activity

Public title

Study of Portulaca Oleracea and Cumin effects on serum paraoxanase 1 activity

Purpose

Treatment

Inclusion/Exclusion criteria

Inclusion criteria: patients with low density lipoprotein cholesterol (LDL-C) over than 120 mg/dl. Exclusion criteria: patients with metabolic disease such as diabetes mellitus, renal diseases and individuals taking lipid lowering or anti hypertension drug.

Age

From **35 years** old to **75 years** old

Gender

Both

Phase

2

Groups that have been masked

No information

Sample size

Target sample size: 90

Randomization (investigator's opinion)

Randomized

Randomization description

Blinding (investigator's opinion)

Not blinded

Blinding description

Placebo

Not used

Assignment

Parallel

Other design features

Secondary Ids

empty

Ethics committees

1

Ethics committee

Name of ethics committee

Shahrekord university of medical sciences

Street address

Kashani street

City

Shahrecord

Postal code

Approval date

2009-02-20, 1387/12/02

Ethics committee reference number

87-10-1

Health conditions studied

1

Description of health condition studied

Metabolic disorders

ICD-10 code

E78

ICD-10 code description

Disorders of lipoprotein metabolism and other lipidaemias

Primary outcomes

1

Description

Paraxonase1

Timepoint

before and after 45 days of intervention

Method of measurement

blood sample

Secondary outcomes

1

Description

Apo A1

Timepoint

before and after 45 days of intervention

Method of measurement

blood sample

2

Description

Apo B

Timepoint

before and after 45 days of intervention

Method of measurement

blood sample

3

Description

Cholesterol

Timepoint

before and after 45 days of intervention

Method of measurement

blood sample

4

Description

Triglyceride

Timepoint

before and after 45 days of intervention

Method of measurement

blood sample

5

Description

HDL

Timepoint

before and after 45 days of intervention

Method of measurement

blood sample

6

Description

LDL

Timepoint

before and after 45 days of intervention

Method of measurement

blood sample

7

Description

OX LDL

Timepoint

before and after 45 days of intervention

Method of measurement

blood sample

Intervention groups

1

Description

Purslane , Almost 50 grams/day of fresh leaves and stems of Purslane, 45 days

Category

Treatment - Other

2

Description

Cumin extract, 1ml of 2% based on manufacturing recommendation , 45 days

Category

Treatment - Drugs

3

Description

Lovastatin, 20mgs/day oral, 45 days

Category

Treatment - Drugs

Recruitment centers

1

Recruitment center

Name of recruitment center

Shahrekord internal clinic, No. 2

Full name of responsible person

Street address

City

Shahrekord

Sponsors / Funding sources

1

Sponsor

Name of organization / entity

Shahrekord university of medical sciences

Full name of responsible person

Dr Mahmood Mobasheri

Street address

Kashani street

City

Shahrekord

Grant name

Grant code / Reference number

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

Yes

Title of funding source

Shahrekord university of medical sciences

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

Shahrekord university of medical sciences

Full name of responsible person

Beheshteh Amiri

Position

nursing

Other areas of specialty/work

Street address

Cellular and molecular research center, Faculty of medicine, Rahmatieh

City

Shahrekord

Postal code

Phone

+98 38 1334 6692

Fax

Email

e_farrokhi_k@yahoo.com

Web page address

Person responsible for scientific inquiries

Contact

Name of organization / entity

Shahrekord university of medical sciences

Full name of responsible person

Dr keihan Ghatreh Samani

Position

PhD in Biochemistry

Other areas of specialty/work

Street address

Biochemistry department, Faculty of medicine, Rahmatieh

City

Shahrekord

Postal code

Phone

+98 38 1333 5654

Fax

Email

kgsamani@yahoo.com

Web page address

Person responsible for updating data

Contact

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