

Clinical Trial Protocol

Iranian Registry of Clinical Trials

11 Jul 2026

Effect of L - carnitine on lipid profile in hyperlipidemic patients referred Hospital Clinic Baqiyatallah

Protocol summary

Summary

The purpose of this study is to evaluate the effect of L-carnitine in 60 hyperlipidemic patients referred to Baqiyatallah special clinic. Diagnosis of hyperlipidemia is based on specialist opinion (cholesterol \geq 200 mg per dL, triglycerides \geq 200 mg per deciliter; aged between 20 to 65 years). Exclusion criteria: risk hypothyroidism, menopause and pregnancy, smoking, drugs and alcohol, inflammatory diseases, such as diabetes, cancer, based on random assignment to intervention and control groups divided. The intervention group received 2 grams L - Carnitine supplement (Company C, followed by Karen Canada) pill daily for 8 weeks (according to studies) and 2 grams placebo pill daily for placebo group. TG, Total Cholesterol, HDL and LDL was calculated before and after the intervention.

General information

Acronym

IRCT registration information

IRCT registration number: **IRCT2014011411920N2**

Registration date: **2014-02-11, 1392/11/22**

Registration timing: **registered_while_recruiting**

Last update:

Update count: **0**

Registration date

2014-02-11, 1392/11/22

Registrant information

Name

Sepideh Abbaszadeh

Name of organization / entity

Baqiyatallah University of Medical Sciences

Country

Iran (Islamic Republic of)

Phone

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Recruitment status

Recruitment complete

Funding source

Baqiyatallah University of Medical Sciences

Expected recruitment start date

2014-01-30, 1392/11/10

Expected recruitment end date

2014-06-22, 1393/04/01

Actual recruitment start date

empty

Actual recruitment end date

empty

Trial completion date

empty

Scientific title

Effect of L - carnitine on lipid profile in hyperlipidemic patients referred Hospital Clinic Baqiyatallah

Public title

Effect of L - carnitine in patients with hyperlipidemia

Purpose

Treatment

Inclusion/Exclusion criteria

Inclusion criteria: risk of hyperlipidemia in the diagnosis and referral specialist measure cholesterol \geq 200 mg dL; triglycerides \geq 200 mg per deciliter; aged between 20 and 65 years; Gender (Female and Male) Exclusion criteria: hypothyroidism; Pregnancy & menopause; smoking; drugs and alcohol; inflammatory diseases; such as diabetes; cancer; chronic liver disease (Fatty Liver, Cirrhosis of the liver, hepatitis, etc.) and use of Anti-seizure drugs; anti-supplements oxidant.

Age

From **20 years** old to **65 years** old

Gender

Both

Phase

N/A

Groups that have been masked

No information

Sample size

Target sample size: 60

Randomization (investigator's opinion)

Randomized

Randomization description**Blinding (investigator's opinion)**

Double blinded

Blinding description**Placebo**

Used

Assignment

Parallel

Other design features**Secondary Ids**

empty

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

Baqiyatallah University of Medical Sciences

Street address

Baqiyatallah University of Medical Science, South sheikh-Bahaei street, Mollasadra Avenue, Tehran, Iran

City

Tehran

Postal code**Approval date**

2013-11-11, 1392/08/20

Ethics committee reference number

340/3/6434/س

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

Hyperlipidemia

ICD-10 code

E78.5

ICD-10 code description

Hyperlipidemia

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

Lipid Profile (Tg,Total Chol, LDL-C, HDL-C)

Timepoint

Baseline and the end of the eighth week

Method of measurement

Their past record of blood test with automated analyzers

Secondary outcomes

empty

Intervention groups**1****Description**

The intervention group completed 8 weeks of daily 2 g L - Carnitine (Company C, followed by Karen Canada) pill of 1000 mg Oral form

Category

Treatment - Drugs

2**Description**

and placebo groups completed 8 weeks 2g daily Oral form use.

Category

Placebo

Recruitment centers**1****Recruitment center****Name of recruitment center**

Baqiyatallah University of Medical Sciences

Full name of responsible person

Shahin Vahabi

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Sponsors / Funding sources**1****Sponsor****Name of organization / entity**

Baqiyatallah University of Medical Sciences

Full name of responsible person

Dr Gashtasb Aghasinejad

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Grant name**Grant code / Reference number****Is the source of funding the same sponsor organization/entity?**

Yes

Title of funding source

Baqiyatallah 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

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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