

Clinical Trial Protocol

Iranian Registry of Clinical Trials

10 Jun 2026

Hypolipidemic effects of Nigella Sativa L. Seeds oil in healthy volunteers: a randomized, double-blind, placebo-controlled clinical trial

Protocol summary

Summary

Background: Several formulations of the Nigella sativa L. seeds (black seed) have been used in traditional medicine for treatment and prevention of a wide range of diseases including hyperlipidemia. But blood lipid lowering effects of its oil in clinical study is of an interest. Objective: The present study was undertaken to explore the possible blood lipid lowering effects of the black seed oil on healthy volunteers. Methods: A randomized clinical trial was conducted in 70 healthy volunteers referring to Bagiatallah hospital. The subjects were randomly selected and enrolled in to two groups of 35 each. One group received 2.5 ml black seed oil and the other group received similarly 2.5 ml mineral oil two times a day. The fasting blood Triglyceride, Cholesterol, HDL, LDL, Glucose, ALT, AST, BUN, Cr, and HbA1C were determined at the baseline and after 8 weeks. Results: Results showed that significant decrease in fasting blood cholesterol, LDL, Triglyceride, glucose and HbA1c levels in black seed oil treated volunteers as compared to placebo group at the end of the study. No notable liver, kidney and gastrointestinal side effects were observed in these two groups side effects were observed in these two groups. Conclusion: Administration of 5 ml black seed oil daily to healthy volunteers for 8 weeks had beneficial effects on improving lipid profile without any adverse effects.

General information

Acronym

IRCT registration information

IRCT registration number: **IRCT201112271157N5**
Registration date: **2012-01-05, 1390/10/15**
Registration timing: **retrospective**

Last update:
Update count: **0**

Registration date

2012-01-05, 1390/10/15

Registrant information

Name

Hasan Fallah Huseini

Name of organization / entity

Institute of Medicinal Plants

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Iran (Islamic Republic of)

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Recruitment status

Recruitment complete

Funding source

Institute of Medicinal Plants, ACECR and Baqiyatallah
University of Medical Sciences

Expected recruitment start date

2011-04-21, 1390/02/01

Expected recruitment end date

2011-05-22, 1390/03/01

Actual recruitment start date

empty

Actual recruitment end date

empty

Trial completion date

empty

Scientific title

Hypolipidemic effects of Nigella Sativa L. Seeds oil in healthy volunteers: a randomized, double-blind, placebo-controlled clinical trial

Public title

Hypolipidemic effects of Nigella Sativa L. seeds oil in healthy volunteers

Purpose

Health service research

Inclusion/Exclusion criteria

Inclusion criteria: healthy Iranian volunteers aged between 40 to 60 years; with fasting blood cholesterol 200 to 300 mg/dl; fasting blood glucose 80 to 120 mg/dl and weight 55 to 85 kg

Age

From **25 years** old to **60 years** old

Gender

Both

Phase

3

Groups that have been masked

No information

Sample size

Target sample size: **70**

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 Sciences, Tehran, Iran

City

Tehran

Postal code

1484958693

Approval date

2010-12-12, 1389/09/21

Ethics committee reference number

22-340/s

Health conditions studied

1

Description of health condition studied

Healthy Volunteers

ICD-10 code

Z01

ICD-10 code description

Other special examinations and investigations of persons without complaint or reported diagnosis

Primary outcomes

1

Description

Cholesterol

Timepoint

At starting of the study and after 8 weeks

Method of measurement

The fasting blood cholesterol level determined in laboratory

Secondary outcomes

1

Description

SGPT

Timepoint

At starting of the study and after 8 weeks

Method of measurement

The fasting blood SGPT level determined in laboratory

2

Description

LDL

Timepoint

At starting of the study and after 8 weeks

Method of measurement

The fasting blood LDL level determined in laboratory

3

Description

Hba1c

Timepoint

At starting of the study and after 8 weeks

Method of measurement

The fasting blood Hba1c level determined in laboratory

4

Description

Glucose

Timepoint

At starting of the study and after 8 weeks

Method of measurement

The fasting blood Glucose level determined in laboratory

5

Description

Creatinine

Timepoint

At starting of the study and after 8 weeks

Method of measurement

The fasting blood Creatinine level determined in laboratory

6

Description

SGOT
Timepoint
At starting of the study and after 8 weeks
Method of measurement
The fasting blood SGOT level determined in laboratory

7

Description
BUN
Timepoint
At starting of the study and after 8 weeks
Method of measurement
The fasting blood BUN level determined in laboratory

8

Description
Triglyceride
Timepoint
At starting of the study and after 8 weeks
Method of measurement
determined in fasting blood sample in laboratory

9

Description
Hdl
Timepoint
At starting of the study and after 8 weeks
Method of measurement
determined in fasting blood sample in laboratory

Intervention groups

1

Description
2.5 ml black seed oil two times a day after meal for 8 weeks
Category
Treatment - Drugs

2

Description
2.5 ml mineral oil two times a day after meal for 8 weeks
Category
Treatment - Drugs

Recruitment centers

1

Recruitment center
Name of recruitment center
Gastroentology and Liver diseases Research Center,
Baqiyatallah University of Medical Sciences
Full name of responsible person
Mohtashami Reza
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Sponsors / Funding sources

1

Sponsor
Name of organization / entity
Institute of Medicinal Plants, ACECR
Full name of responsible person
Hassan ali Nqdi badi
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Tehran-Qazwain Highway Research City Kavosh,
Jahad daneshgahi research society Kraj
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Kraj
Grant name
روغن سیاه دانه: 84 / 1389/9/10 /س
Grant code / Reference number
1142-33
Is the source of funding the same sponsor organization/entity?
Yes
Title of funding source
Institute of Medicinal Plants, ACECR
Proportion provided by this source
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

2

Sponsor
Name of organization / entity
Baqiyatallah University of Medical Sciences
Full name of responsible person
Reza Mohtashami
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Baqiyatallah University of Medical Sciences, Tebo-o-
Din Center, Vanak squer, Molasadra Ave, Tehran, Iran
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Grant name
456/1/1389
Grant code / Reference number
111-306
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
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

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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