

# Clinical Trial Protocol

## Iranian Registry of Clinical Trials

27 Jun 2026

### Comparison of the effects of Nigella sativa seed oil mixed with honey and placebo on the hepatic sonography and blood biochemical parameters of non-alcoholic fatty liver disease patients

#### Protocol summary

##### Study aim

Comparison of the effectiveness of Nigella sativa seed oil mixed with honey and placebo in the treatment of non-alcoholic fatty liver disease patients

##### Design

Randomized; block randomization with computer generated random numbers table and sequentially numbered containers each representing a block consisting of ten patients are used for the treatment assignments; double-blind; controlled with placebo; two group parallel design; single-center; phase 3 of clinical trial. Study population: patients with non-alcoholic fatty liver disease. Sample size: 120 patients.

##### Settings and conduct

The trial setting: Baqiyatallah Hospital. Protocol: sixty patients use 2.5 mL of Nigella sativa seed oil and 60 patients use 2.5 mL placebo orally every 12 hours for 3 months. Before intervention and 3 months after intervention the groups' liver sonography and blood levels of ALT, AST, LDL-C, HDL-C and triglyceride are compared with each other. Blinding: three different persons generate the random allocation sequence, enroll the participants and assign them to intervention groups. These persons, care providers, outcome assessors, data analyzer, investigators and participants are blinded to assignment to the study groups.

##### Participants/Inclusion and exclusion criteria

Inclusion criteria: patients whose hepatic sonography shows they have fatty liver disease; age of 20 to 70 years. Exclusion criteria: Child-Pugh score above 7; hepatic disease other than non-alcoholic fatty liver disease; pregnant women; women planning pregnancy; lactating women.

##### Intervention groups

Sixty patients use 2.5 mL of Nigella sativa seed oil and 60 patients use 2.5 mL of placebo orally every 12 hours for 3 months.

#### Main outcome variables

Primary outcome variables: fatty liver grade, blood level of ALT, blood level of AST, blood level of LDL-C, blood level of triglyceride. Secondary outcome variable: blood level of HDL-C.

#### General information

##### Reason for update

##### Acronym

##### IRCT registration information

IRCT registration number: **IRCT20090804002288N13**

Registration date: **2018-06-06, 1397/03/16**

Registration timing: **retrospective**

Last update: **2018-06-06, 1397/03/16**

Update count: **0**

##### Registration date

2018-06-06, 1397/03/16

##### Registrant information

##### Name

Saeed Kianbakht

##### Name of organization / entity

Iranian Academic Center for Education, Culture and Research Institute of Medicinal Plants

##### Country

Iran (Islamic Republic of)

##### Phone

+98 26147640109

##### Email address

kianbakht@imp.ac.ir

##### Recruitment status

**Recruitment complete**

##### Funding source

##### Expected recruitment start date

2017-04-04, 1396/01/15

**Expected recruitment end date**

2018-04-20, 1397/01/31

**Actual recruitment start date**

2017-04-04, 1396/01/15

**Actual recruitment end date**

2018-04-20, 1397/01/31

**Trial completion date**

empty

**Scientific title**

Comparison of the effects of Nigella sativa seed oil mixed with honey and placebo on the hepatic sonography and blood biochemical parameters of non-alcoholic fatty liver disease patients

**Public title**

Effect of Nigella sativa seed oil in treatment of non-alcoholic fatty liver disease

**Purpose**

Treatment

**Inclusion/Exclusion criteria****Inclusion criteria:**

Patients whose hepatic sonography shows they have fatty liver disease Age of 20 to 70 years

**Exclusion criteria:**

Child-Pugh score above 7 Hepatic disease other than non-alcoholic fatty liver disease Pregnant women Women planning pregnancy Lactating women

**Age**

From **20 years** old to **70 years** old

**Gender**

Both

**Phase**

N/A

**Groups that have been masked**

- Participant
- Care provider
- Investigator
- Outcome assessor
- Data analyser

**Sample size**

Target sample size: **120**

Actual sample size reached: **120**

**Randomization (investigator's opinion)**

Randomized

**Randomization description**

Block randomization with computer generated random numbers table and sequentially numbered containers each representing a block consisting of ten patients are used for the treatment assignments.

**Blinding (investigator's opinion)**

Double blinded

**Blinding description**

Three different persons generate the random allocation sequence, enroll the participants and assign them to intervention groups. These persons, care providers, outcome assessors, data analyzer, investigators and participants are blinded to assignment to the Nigella sativa seed oil and placebo groups.

**Placebo**

Used

**Assignment**

Parallel

**Other design features****Secondary Ids**

empty

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

Ethics committee of Baqiyatallah University of Medical Sciences

**Street address**

Molla Sadra Street, Vanak Square

**City**

Tehran

**Province**

Tehran

**Postal code**

1435915371

**Approval date**

2015-11-01, 1394/08/10

**Ethics committee reference number**

IR.BMSU.REC.1394.149

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

Non-alcoholic fatty liver disease

**ICD-10 code**

K76.0

**ICD-10 code description**

Fatty (change of) liver, not elsewhere classified

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

Fatty liver grade

**Timepoint**

Before intervention and 3 month پس از intervention

**Method of measurement**

Liver sonography

**2****Description**

Blood level of ALT

**Timepoint**

Before intervention and 3 months after intervention

**Method of measurement**

Medical laboratory (kit)

### 3

**Description**

Blood level of AST

**Timepoint**

Before intervention and 3 months after intervention

**Method of measurement**

Medical laboratory (kit)

### 4

**Description**

Blood level of LDL-C

**Timepoint**

Before intervention and 3 months after intervention

**Method of measurement**

Medical laboratory (kit)

### 5

**Description**

Blood level of triglyceride

**Timepoint**

Before intervention and 3 months after intervention

**Method of measurement**

Medical laboratory (kit)

## Secondary outcomes

### 1

**Description**

Blood level of HDL-C

**Timepoint**

Before intervention and 3 month after intervention

**Method of measurement**

Medical laboratory (kit)

## Intervention groups

### 1

**Description**

Intervention group: sixty patients use 2.5 mL of Nigella sativa seed oil orally every 12 hours for 3 months.

**Category**

Treatment - Drugs

### 2

**Description**

Control group: sixty patients use 2.5 mL of placebo orally every 12 hours for 3 months.

**Category**

Placebo

## Recruitment centers

### 1

**Recruitment center**

Name of recruitment center

Baqiyatallah Hospital

**Full name of responsible person**

Dr. Reza Mohtashami

**Street address**

Molla Sadra Street, Vanak Square

**City**

Tehran

**Province**

Tehran

**Postal code**

1435915371

**Phone**

+98 21 8805 0435

**Email**

reza\_mohtashami1979@yahoo.com

## Sponsors / Funding sources

### 1

**Sponsor****Name of organization / entity**

Iranian academic center for education culture and research

**Full name of responsible person**

Dr. Reza Hajiaghaee

**Street address**

ACECR Complex, Supa Boulevard, Poleh Kordan

**City**

Karaj

**Province**

Alborz

**Postal code**

3365166571

**Phone**

+98 26 3476 4010

**Email**

hajiaghaee@imp.ac.ir

**Web page address**

<http://www.imp.ac.ir>

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

Yes

**Title of funding source**

Iranian academic center for education culture and research

**Proportion provided by this source**

100

**Public or private sector**

Public

**Domestic or foreign origin**

Domestic

**Category of foreign source of funding**

*empty*

**Country of origin****Type of organization providing the funding**

Academic

## Person responsible for general inquiries

### Contact

**Name of organization / entity**

Iranian academic center for education culture and research

**Full name of responsible person**

Dr. Saeed Kianbakht

**Position**

Assistant professor

**Latest degree**

Ph.D.

**Other areas of specialty/work**

Pharmacology

**Street address**

Institute of Medicinal Plants, ACECR Complex, Supa Boulevard, Poleh Kordan

**City**

Karaj

**Province**

Alborz

**Postal code**

3365166571

**Phone**

+98 26 3476 4010

**Email**

kianbakht@imp.ac.ir

**Web page address**

<http://www.imp.ac.ir>

## Person responsible for scientific inquiries

### Contact

**Name of organization / entity**

Iranian academic center for education culture and research

**Full name of responsible person**

Dr. Saeed Kianbakht

**Position**

Assistant professor

**Latest degree**

Ph.D.

**Other areas of specialty/work**

Pharmacology

**Street address**

Institute of Medicinal Plants, ACECR Complex, Supa Boulevard, Poleh Kordan

**City**

Karaj

**Province**

Alborz

**Postal code**

3365166571

**Phone**

+98 26 3476 4010

**Email**

kianbakht@imp.ac.ir

**Web page address**

<http://www.imp.ac.ir>

## Person responsible for updating data

### Contact

**Name of organization / entity**

Iranian academic center for education culture and research

**Full name of responsible person**

Dr. Saeed Kianbakht

**Position**

Assistant professor

**Latest degree**

Ph.D.

**Other areas of specialty/work**

Pharmacology

**Street address**

Institute of Medicinal Plants, ACECR Complex, Supa Boulevard, Poleh Kordan

**City**

Karaj

**Province**

Alborz

**Postal code**

3365166571

**Phone**

+98 26 3476 4010

**Email**

kianbakht@imp.ac.ir

**Web page address**

<http://www.imp.ac.ir>

## Sharing plan

**Deidentified Individual Participant Data Set (IPD)**

Yes - There is a plan to make this available

**Study Protocol**

Yes - There is a plan to make this available

**Statistical Analysis Plan**

Yes - There is a plan to make this available

**Informed Consent Form**

No - There is not a plan to make this available

**Clinical Study Report**

Yes - There is a plan to make this available

**Analytic Code**

No - There is not a plan to make this available

**Data Dictionary**

No - There is not a plan to make this available

**Title and more details about the data/document**

All collected deidentified IPD are to be shared.

**When the data will become available and for how long**

Access period starting 6 months after publication for 1 year.

**To whom data/document is available**

Researchers working in university and scientific and industrial institutes.

**Under which criteria data/document could be used**

Only statistical analyses mentioned in the article resulting from the study for personal knowledge are permitted. Written request for access to data and documents and its aim must be certified by the highest ranking official of the work place of the requester.

**From where data/document is obtainable**

Dr. Saeed Kianbakht with the address ACECR Institute of Medicinal plants, P.O. Box: 31375-369, Iran.

**What processes are involved for a request to access data/document**

Written request should be sent to the address Dr. Saeed

Kianbakht, ACECR Institute of Medicinal Plants, P.O. Box: 31375-369, Iran. The documents or data file will be sent to the requester by email.

**Comments**