

Clinical Trial Protocol

Iranian Registry of Clinical Trials

10 Jun 2026

Evaluating the effect of black garlic on liver steatosis and enzymes, lipid profile, insulin resistance, and anthropometric measurements in patients with non-alcoholic fatty liver disease

Protocol summary

Study aim

Determining the effect of black garlic on liver steatosis and enzymes, lipid profile, insulin resistance, and anthropometric measurements in patients with non-alcoholic fatty liver disease

Design

Parallel randomized controlled trial

Settings and conduct

The present study aiming to evaluate effect of black garlic on liver steatosis and enzymes, lipid profile, insulin resistance, and anthropometric measurements in patients with non-alcoholic fatty liver disease. This study will take place in Imam Khomeini Hospital in Urmia and it is a parallel randomized controlled clinical trial. 110 patients will be randomly assigned to 2 groups.

Participants/Inclusion and exclusion criteria

Inclusion criteria: Patients with non-alcoholic fatty liver disease more than 18 years from both sexes Patients with non-alcoholic fatty liver disease with a BMI less than 30; patients with non-alcoholic fatty liver disease diagnosed through ultrasound and referred to Imam Hospital Gastroenterology Clinic. Exclusion criteria: Special diets; pregnancy or lactation during the study; viral hepatitis; alcohol consumption or history of alcohol consumption; non-compliance to our recommendations regarding Black garlic consumption; untreated hypothyroidism; renal diseases; taking drugs that increase liver enzymes; alpha antitrypsin deficiency; gastrointestinal disease such as celiac disease; menopause; diabetes mellitus; herbal therapy; heart failure and bone disease.

Intervention groups

In the intervention group, in addition to the usual treatment, will be recommended to consume 6 grams of black garlic in total per day before morning and evening meal. In addition to the usual fatty liver treatment, the control group was advised to follow FAO (Food and

Agriculture Organization) recommendations by the researcher.

Main outcome variables

Liver steatosis; Insulin resistance; Liver enzyme; Lipid profile; Anthropometric measurements

General information

Reason for update

Acronym

IRCT registration information

IRCT registration number: **IRCT20201204049599N6**

Registration date: **2022-10-26, 1401/08/04**

Registration timing: **prospective**

Last update: **2022-10-26, 1401/08/04**

Update count: **0**

Registration date

2022-10-26, 1401/08/04

Registrant information

Name

Mohammad Reza Pashaei

Name of organization / entity

Country

Iran (Islamic Republic of)

Phone

+98 44 3198 8001

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Recruitment status

Recruitment complete

Funding source

Expected recruitment start date

2022-11-05, 1401/08/14

Expected recruitment end date

2023-03-16, 1401/12/25

Actual recruitment start date

empty

Actual recruitment end date

empty

Trial completion date

empty

Scientific title

Evaluating the effect of black garlic on liver steatosis and enzymes, lipid profile, insulin resistance, and anthropometric measurements in patients with non-alcoholic fatty liver disease

Public title

Effect of Black garlic consumption on treatment of non-alcoholic fatty liver disease

Purpose

Treatment

Inclusion/Exclusion criteria**Inclusion criteria:**

Age older than 18 years old from both sexes BMI more than 30 Patients with non-alcoholic fatty liver disease diagnosed through ultrasound and referred to Imam Hospital Gastroenterology Clinic

Exclusion criteria:

Special diets Pregnancy and lactation during the study Viral hepatitis Alcohol consumption or history of alcohol consumption Non-compliance to our recommendations regarding Black garlic consumption Untreated hypothyroidism Renal diseases Taking drugs that increase liver enzymes Alpha antitrypsin deficiency Gastrointestinal disease such as celiac disease Menopause Diabetes mellitus Herbal therapy Heart failure and bone disease

Age

From **18 years** old

Gender

Both

Phase

N/A

Groups that have been masked

No information

Sample size

Target sample size: **56**

Randomization (investigator's opinion)

Randomized

Randomization description

The method used for randomization is Stratified Randomization. In this method, to eliminate the heterogeneity in age, gender and severity of steatosis variables, before applying random allocation, the samples will be classified based on these variables and then they will be randomly allocated into intervention and control groups.

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**

Research Ethics Committee of Urmia University of Medical Sciences - Imam Khomeini University Hospi

Street address

Imam Khomeini Hospital; Ershad Blvd., Urmia, Iran

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Urmia

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West Azarbaijan

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5715789397

Approval date

2022-08-30, 1401/06/08

Ethics committee reference number

IR.UMSU.HIMAM.REC.1401.041

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**

Weight

Timepoint

At weeks 0, 12

Method of measurement

Scale, kg

2**Description**

Body mass index

Timepoint

At weeks 0, 12

Method of measurement

Weight(kg)/[height(m)]² , kg/m²

3**Description**

Gamma Glutamyl transferase

Timepoint

At weeks 0,12

Method of measurement
Enzymatic method , IU/ Lit

4

Description
Alkaline Phosphatase
Timepoint
At weeks 0, 12
Method of measurement
Enzymatic method , IU/ Lit

5

Description
Aspartate transaminase
Timepoint
At weeks 0, 12
Method of measurement
Enzymatic method, IU/Lit

6

Description
Alanine Aminotransferase
Timepoint
At weeks 0, 12
Method of measurement
Enzymatic method, IU/ Lit

7

Description
Fasting blood sugar
Timepoint
At weeks 0, 12
Method of measurement
Enzymatic method, IU/ Lit

8

Description
Serum insulin
Timepoint
At weeks 0, 12
Method of measurement
Radioimmunoassay

9

Description
Waist circumference
Timepoint
At weeks 0, 12
Method of measurement
Cm

10

Description
Hepatic steatosis
Timepoint
Before intervention and three months after intervention
Method of measurement

Ultrasonography

11

Description
HDL cholesterol
Timepoint
At weeks 0, 12
Method of measurement
Enzymatic method, mg/dl

12

Description
Triglyceride
Timepoint
At weeks 0, 12
Method of measurement
Enzymatic method, mg/dl

13

Description
LDL cholesterol
Timepoint
At weeks 0, 12
Method of measurement
Enzymatic method, mg/dl

14

Description
Total cholesterol
Timepoint
At weeks 0, 12
Method of measurement
Enzymatic method, mg/dl

15

Description
HOMA-IR
Timepoint
At weeks 0, 12
Method of measurement
Formula

16

Description
Height
Timepoint
Before intervention
Method of measurement
Stadiometer, mm

Secondary outcomes

empty

Intervention groups

1

Description

Intervention group: In the intervention group, in addition to the usual treatment, will be recommended to consume 6 grams of black garlic in total per day before morning and evening meal for three months.

Category

Treatment - Other

2

Description

Control group: In addition to the usual fatty liver treatment, the control group was advised to follow FAO (Food and Agriculture Organization) recommendations by the researcher for three months.

Category

Treatment - Other

Recruitment centers

1

Recruitment center

Name of recruitment center

Imam Khomeini Teaching Hospital, Urmia

Full name of responsible person

Mohammad Reza Pashaei

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Imam Khomeini Hospital; Ershad Blvd., Urmia, Iran

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Sponsors / Funding sources

1

Sponsor

Name of organization / entity

Oroumia University of Medical Sciences

Full name of responsible person

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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

Oroumia University of Medical Sciences

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

Oroumia University of Medical Sciences

Full name of responsible person

Mohammad Reza Pashaei

Position

Assistant professor

Latest degree

Subspecialist

Other areas of specialty/work

Internal Medicine

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Latest degree

Subspecialist

Other areas of specialty/work

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Person responsible for updating data

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Sharing plan

Deidentified Individual Participant Data Set (IPD)

Undecided - It is not yet known if there will be a plan to make this available

Study Protocol

Undecided - It is not yet known if there will be a plan to make this available

Statistical Analysis Plan

Undecided - It is not yet known if there will be a plan to make this available

Informed Consent Form

Undecided - It is not yet known if there will be a plan to make this available

Clinical Study Report

Undecided - It is not yet known if there will be a plan to make this available

Analytic Code

Undecided - It is not yet known if there will be a plan to make this available

Data Dictionary

Undecided - It is not yet known if there will be a plan to make this available