

Clinical Trial Protocol

Iranian Registry of Clinical Trials

08 Jun 2026

Evaluation of the effects of rice bran oil administration on metabolic syndrome components, inflammatory condition and Apo A1 to Apo B100 ratio in adult patients with metabolic syndrome: An Open Label Randomized Controlled Trial

Protocol summary

Study aim

Determining the effects of rice bran oil administration on the indicators of metabolic syndrome, inflammatory status and the ratio of apolipoproteins A1 to B100 in adult patients with metabolic syndrome

Design

Clinical trial with control group, with parallel groups, without blinding, randomized, phase 2 on 50 patients to allocate consumption to the subjects will use the randomized block method. The website <http://www.randomization.com> will also be used for randomization

Settings and conduct

Fifty patients with metabolic syndrome referred to the outpatient clinic of Dr. Heshmat Heart Training Center will be admitted with personal consent after completing the informed consent form, taking into account the inclusion and exclusion criteria. Patients in two groups of 25 will be treated with rice bran oil (2 tablespoons daily) and standard diet or standard diet with 30 grams or 2 tablespoons per day of sunflower oil (as a control group).

Participants/Inclusion and exclusion criteria

Patients with an age range of 20 to 70 years, Patients with metabolic syndrome, Do not use vitamin and mineral supplements, antioxidants, fiber supplements, omega 3, No history of kidney disease, kidney stones, gastrointestinal diseases, gallstones, and autoimmune diseases, Current consumption of alcohol

Intervention groups

Intervention group: Patients are asked to consume 30 grams of rice bran oil, which is equivalent to 2 tablespoons of bran oil prepared by Giltaz Company, along with lunch. Control group: Patients are asked to consume 30 grams of sunflower oil, which is equivalent to 2 tablespoons of oil prepared from authorized stores, along with lunch.

Main outcome variables

Cardiovascular risk factors

General information

Reason for update

Acronym

IRCT registration information

IRCT registration number: **IRCT20180205038626N10**

Registration date: **2021-11-25, 1400/09/04**

Registration timing: **registered_while_recruiting**

Last update: **2021-11-25, 1400/09/04**

Update count: **0**

Registration date

2021-11-25, 1400/09/04

Registrant information

Name

Zahra Ahmadnia

Name of organization / entity

Country

Iran (Islamic Republic of)

Phone

+98 13 3361 8177

Email address

zahmadnia@gums.ac.ir

Recruitment status

Recruitment complete

Funding source

Expected recruitment start date

2021-11-22, 1400/09/01

Expected recruitment end date

2022-11-22, 1401/09/01

Actual recruitment start date

empty

Actual recruitment end date
empty

Trial completion date
empty

Scientific title
Evaluation of the effects of rice bran oil administration on metabolic syndrome components, inflammatory condition and Apo A1 to Apo B100 ratio in adult patients with metabolic syndrome: An Open Label Randomized Controlled Trial

Public title
Evaluation of the effects of rice bran oil administration on metabolic syndrome components, inflammatory condition and Apo A1 to Apo B100 ratio in adult patients with metabolic syndrome

Purpose
Prevention

Inclusion/Exclusion criteria
Inclusion criteria:
Patients with an age range of 20 to 70 years Patients with metabolic syndrome Do not use vitamin and mineral supplements, antioxidants, fiber supplements, omega 3 No history of kidney disease, kidney stones, gastrointestinal diseases, gallstones, and autoimmune diseases No current consumption of alcohol
Exclusion criteria:
Changes in the patient's treatment plan during the study Changing the type of effective drugs used factors studied Reluctance to continue the study or to cause any dissatisfaction with the taste of the oil or to participate in the study

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

Gender
Both

Phase
2

Groups that have been masked
No information

Sample size
Target sample size: **50**

Randomization (investigator's opinion)
Randomized

Randomization description
Sampling and randomization The clinical trial study of two parallel groups of open label will be classified by block random sampling method. At first, participants were classified into two classes according to age (20 to 45 years and between 45 to 70 years) and then each person was randomly assigned to the intervention or control group using 1: 4 random blocks. Took. In this method, each group will be assigned one of the letters A or B. The website will also be used for randomization. The list of codes obtained from this website will be provided to the researchers, and each referring patient who met the inclusion criteria and did not meet the inclusion criteria and was willing to participate in the study, first entered the desired age group and based on The assigned code A or B enters the design. For

concealment, in this study, random allocation concealment, which is the method used to execute a random sequence on the study participants, will be used in such a way that the assigned group is not known before the individual is assigned. In this way, using opaque envelopes sealed with a random sequence, in this method, each of the random sequences created is recorded on a card and the cards are placed inside the envelopes in order. In order to maintain a random sequence, the envelopes are numbered in the same way on the outer surface. Finally, the lids of the letter envelopes are glued and placed inside a box, respectively. At the beginning of the registration of participants, based on the order of entry of eligible participants into the study, one of the envelopes of the letter is opened in order and the assigned group of the participant is revealed.

Blinding (investigator's opinion)

Not blinded

Blinding description

Placebo

Used

Assignment

Parallel

Other design features

Secondary Ids

empty

Ethics committees

1

Ethics committee

Name of ethics committee

Ethics committee of Guilan University of Medical Sciences

Street address

Technology & Research Vice-chancellor of University; Shahid Siadati St; Namjoo St., Rasht

City

Rasht

Province

Guilan

Postal code

41446-66949

Approval date

2021-11-10, 1400/08/19

Ethics committee reference number

IR.GUMS.REC.1400.389

Health conditions studied

1

Description of health condition studied

Metabolic syndrome

ICD-10 code

E88.81

ICD-10 code description

Metabolic syndrome

Primary outcomes

1

Description

Serum lipid profile levels (triglycerides, total cholesterol, LDL-C and HDL-C)

Timepoint

At the beginning of the study and 8 weeks later

Method of measurement

BT2000 devices

2

Description

Serum levels of Apo A1 and Apo B100

Timepoint

At the beginning of the study and 8 weeks later

Method of measurement

Nefrometric devices

3

Description

Fasting blood sugar

Timepoint

At the beginning of the study and 8 weeks later

Method of measurement

BT2000 devices

4

Description

CRP

Timepoint

At the beginning of the study and 8 weeks later

Method of measurement

ELISA Devices

5

Description

D-dimer

Timepoint

At the beginning of the study and 8 weeks later

Method of measurement

ELISA Device

Secondary outcomes

1

Description

Mean systolic blood pressure

Timepoint

At the beginning of the study and 8 weeks later

Method of measurement

Pressure indicator

2

Description

Mass Index body

Timepoint

At the beginning of the study and 8 weeks later

Method of measurement

Centimeter and scales

3

Description

Waist

Timepoint

At the beginning of the study and 8 weeks later

Method of measurement

Centimeter

4

Description

mean diastolic blood pressure

Timepoint

At the beginning of the study and 8 weeks later

Method of measurement

Pressure indicator

Intervention groups

1

Description

Intervention group: Patients with metabolic syndrome will be included in the study for 8 weeks to receive a standard diet + consumption of 30 grams of rice bran oil per day. Patients in the intervention group are asked to consume 30 grams of rice bran oil, which is equivalent to 2 tablespoons of bran oil prepared by Giltaz Company, along with lunch.

Category

Prevention

2

Description

Control group: Patients in the control group are also asked to consume 30 grams of sunflower oil, which is equivalent to 2 tablespoons of oil prepared from authorized stores, along with lunch. Oils are given to all patients in a graduated dose and they are asked to add 30 grams of it daily in a graduated dose to their salad or cooked food at lunch.

Category

Prevention

Recruitment centers

1

Recruitment center

Name of recruitment center

Heshmat hospital

Full name of responsible person

Zeynab Ghorbani Lohesara

Street address

15 Khordad St., next to the Management and

Planning Organization of Gilan Province, Dr. Heshmat
Heart Training and Treatment Center

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4193955588

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Email

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Sponsors / Funding sources

1

Sponsor

Name of organization / entity

Rasht University of Medical Sciences

Full name of responsible person

Mohammadreza Naghipour

Street address

Technology & Research Vice-chancellor of University;
Shahid Siadati St; Namjoo St., Rasht

City

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41446-66949

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+98 13 3333 5821

Email

naghi@gums.ac.ir

Grant name

Grant code / Reference number

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

Yes

Title of funding source

Rasht 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

Rasht University of Medical Sciences

Full name of responsible person

Zahra Ahmadnia

Position

Nurse

Latest degree

Master

Other areas of specialty/work

Nursery

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Dr. Heshmat Hospital; Bayani St; Mosala Square;
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Person responsible for scientific inquiries

Contact

Name of organization / entity

Rasht University of Medical Sciences

Full name of responsible person

Zeynab Ghorbani lohesara

Position

Assistant Professor

Latest degree

Ph.D.

Other areas of specialty/work

Nutrition

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Person responsible for updating data

Contact

Name of organization / entity

Rasht University of Medical Sciences

Full name of responsible person

Zahra Ahmadnia

Position

NURSE

Latest degree

Master

Other areas of specialty/work

Nursery

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Sharing plan

Deidentified Individual Participant Data Set (IPD)

Yes - There is 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

Title and more details about the data/document

Information on the main outcome

When the data will become available and for how long

Access period starts 6 months after the results are published

To whom data/document is available

Researchers working in academic institutions

Under which criteria data/document could be used

Request to receive unidentifiable personal data or other documents

From where data/document is obtainable

Zeynab Ghorbany Lohesara

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

By email to dr.zeynab.ghorbani.lohesara

Comments