

Clinical Trial Protocol

Iranian Registry of Clinical Trials

09 Jun 2026

The effect of moderately-restricted carbohydrate diet on intestinal microbiota and metabolic parameters in women with metabolic syndrome

Protocol summary

Intestinal microbiome composition, components of metabolic syndrome

Study aim

The effect of moderately-restricted carbohydrate diet on intestinal microbiota and metabolic parameters in women with metabolic syndrome

Design

A clinical trial with a control group, with parallel groups, randomized on 70 patients

Settings and conduct

This study is a two-stage randomized clinical trial study that will be performed on adult women (20-50 years) with metabolic syndrome. The "invitation to participate in the research project" will be used in the Nutrition Clinic of Shariati Hospital, Tehran to reach the mentioned people. People will be randomly placed in a double block in terms of age and body mass index. The control group will be on a normal weight loss diet for 3 months, and the intervention group will be on a diet with moderate carbohydrate intake for 3 months.

Participants/Inclusion and exclusion criteria

Metabolic syndrome based on ATP III criteria, female gender, age 20-50 years, no menopause, overweight and obesity (BMI above 25), not pregnant and lactating, and willing to participate in the study. The study included: pregnancy at any stage of the study, smoking, chronic cardiovascular disease, liver, kidney disease, inflammatory bowel disease such as colitis, thyroid, diabetes, rheumatoid arthritis, cancer, lupus, severe infection, and trauma A history of weight loss surgery, a serious allergy to a particular drug or food, following a specific diet for the 3 months before the study, taking antibiotics and probiotic supplements over the past three months, and taking medications that affect weight.

Intervention groups

People in the intervention group will receive a weight loss diet with a limit of 500 kcal and a moderate carbohydrate intake limitation. People in the control group will receive a normal weight loss diet with a limit of 500 kcal.

Main outcome variables

General information

Reason for update

Acronym

IRCT registration information

IRCT registration number: **IRCT20210307050621N1**

Registration date: **2021-05-12, 1400/02/22**

Registration timing: **prospective**

Last update: **2021-05-12, 1400/02/22**

Update count: **0**

Registration date

2021-05-12, 1400/02/22

Registrant information

Name

Seyed Mohammad Mousavi

Name of organization / entity

Country

Iran (Islamic Republic of)

Phone

+98 21 8895 5742

Email address

smmousavi1993@yahoo.com

Recruitment status

Recruitment complete

Funding source

Expected recruitment start date

2021-09-23, 1400/07/01

Expected recruitment end date

2022-04-21, 1401/02/01

Actual recruitment start date

empty

Actual recruitment end date

empty

Trial completion date

empty

Scientific title

The effect of moderately-restricted carbohydrate diet on intestinal microbiota and metabolic parameters in women with metabolic syndrome

Public title

The effect of moderately-restricted carbohydrate diet on intestinal microbiota

Purpose

Treatment

Inclusion/Exclusion criteria**Inclusion criteria:**

Having metabolic syndrome based on The Adult Treatment Panel guidelines (ATP III) index Age 20 to 50 years Not being menopausal Body mass index greater than or equal to 25 Willingness to participate in the study

Exclusion criteria:

Pregnancy or breastfeeding or deciding to get pregnant shortly Smoking even one cigarette or once using other tobacco such as hookah in the past week or repeated drug use History of the liver, kidney, thyroid, gastrointestinal diseases, diabetes, rheumatoid arthritis, lupus, severe infection, and trauma History of weight loss surgery Having a serious allergy to a particular medicine or food Follow a special diet for 3 months before the study Take any amount of supplements or food products fortified with fins or probiotics during the last 3 months Taking a variety of antibiotics per dose during the 3 months before enrollment Taking drugs that affect weight, appetite, blood pressure, or metabolism of carbohydrates and fats in the last 3 months

Age

From **20 years** old to **50 years** old

Gender

Female

Phase

N/A

Groups that have been masked

No information

Sample size

Target sample size: **70**

Randomization (investigator's opinion)

Randomized

Randomization description

First, subjects will be placed in double blocks in terms of age and body mass index. Then, randomly assigned individuals in each block will be assigned to intervention and control groups. To randomly assign individuals to groups, each person is assigned a code, and these codes are poured into a pot. An out-of-study person is then asked to draw the codes out of the pot using a lottery. The first code will be assigned to the intervention group, the second code to the control group, and so the rest of the people will be randomly assigned to the two 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**

Ethics committee of Tehran University of Medical Sciences

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Faculty of Nutrition and Dietetics, University of Medical Sciences and Health Services, Tehran, Keshavarz Blvd., Naderi St., Hojjatdoost Alley, No. 44

City

Tehran

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Postal code

1416643931

Approval date

2021-04-28, 1400/02/08

Ethics committee reference number

IR.TUMS.MEDICINE.REC.1400.116

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**

Intestinal microbiome composition

Timepoint

Before the intervention and 3 months later

Method of measurement

Polymerase Chain Reaction (PCR) method

2**Description**

Lipid profile (serum triglyceride, total serum cholesterol, low-density lipoprotein, high-density lipoprotein)

Timepoint

Before the intervention and 3 months later

Method of measurement

Enzymatic method

3

Description

Waist circumference

Timepoint

Before the intervention and 3 months later

Method of measurement

Tape meter

4

Description

Fasting blood sugar

Timepoint

Before the intervention and 3 months later

Method of measurement

By glucose oxidase method and using commercial kits

Secondary outcomes

1

Description

Body weight

Timepoint

Before the intervention and 3 months later

Method of measurement

Scale

2

Description

Blood pressure

Timepoint

Before the intervention and 3 months after the intervention

Method of measurement

Sphygmomanometer

3

Description

Height

Timepoint

Before the intervention and 3 months later

Method of measurement

Meter

Intervention groups

1

Description

Intervention group: Regular weight loss diet (daily limit of 500 kcal) and moderate carbohydrate restriction include 45-42% carbohydrates, 40-35% fat, and 17-15% protein. This diet should be consumed daily by participants for 3 months.

Category

Lifestyle

2

Description

Control group: The usual weight loss diet (daily limit of 500 kcal) includes 55-52% carbohydrates, 25-25% fat and 17-15% protein. This diet should be observed daily by participants for 3 months.

Category

Lifestyle

Recruitment centers

1

Recruitment center**Name of recruitment center**

Using "Call to participate in a research project" in various places and in private offices in Tehran

Full name of responsible person

Dr Ahmad Esmailzadeh

Street address

Faculty of Nutrition and Dietetics, No. 44, Hojjatdoost Alley, Naderi St., Keshavarz Blvd., Keshavarz Blvd., Tehran University of Medical Sciences and Health Services

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Sponsors / Funding sources

1

Sponsor**Name of organization / entity**

Tehran 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

Tehran University of Medical Sciences

Proportion provided by this source

60

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**

Tehran University of Medical Sciences

Full name of responsible person

Seyed Mohammad Mousavi

Position

Ph.D Student

Latest degree

Master

Other areas of specialty/work

Nutrition

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Faculty of Nutrition and Dietetics, No. 44, Hojjatdoost Alley, Naderi St., Keshavarz Blvd., Keshavarz Blvd., Tehran University of Medical Sciences and Health Services

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Person responsible for scientific inquiries**Contact****Name of organization / entity**

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Full name of responsible person

Dr Ahmad Esmailzadeh

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Professor of nutrition

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Person responsible for updating data**Contact****Name of organization / entity**

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Full name of responsible person

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Position

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Other areas of specialty/work

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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

Yes - There is a plan to make this available

Statistical Analysis Plan

No - There is not 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

The study protocol will be written and published in the

form of an article. The clinical report of the study will be published in the form of an article.

When the data will become available and for how long

8 months after the end of the study

To whom data/document is available

The information will be made available to the public.

Under which criteria data/document could be used

To use the findings in the clinic or to write other articles, including review articles. In the case of original articles,

researchers will be allowed to do so.

From where data/document is obtainable

Update information via email to the person in charge

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

After receiving the request from the person in charge of updating, the study will be provided to the researcher in consultation with the scientific officer.

Comments