

Clinical Trial Protocol

Iranian Registry of Clinical Trials

19 Jun 2026

The influence of probiotics and Zinc on the serum levels of liver enzymes and lipid profile in patients with non-alcoholic fatty liver disease

Protocol summary

Summary

The aim of this study is assess the influence of probiotics and Zinc on the serum levels of liver enzymes and lipid profile in patients with non-alcoholic fatty liver disease (NAFLD). This is a randomized, triple blind (blinding for patients, doctors and he who analyzed), controlled with placebo and single central study. In this study, 128 patients with NAFLD are studied. Inclusion criteria were Patients with NAFLD and exclusion criteria were autoimmune hepatitis, viral hepatitis, liver metabolic diseases, age below 18 and etc. Patients were randomly (randomized block design) divided into 4 groups of 32 people. The first group receive two probiotic capsules and two zinc capsules, the second group received two probiotic capsules and two zinc concentrate capsules, the third group received two capsules of probiotic placebo and two zinc capsules, and the fourth group received two probiotic capsules and two zinc plaques placebo capsules daily. The intervention time is 3 months. The serum level of ALT, AST, FBS, TG, Total cholesterol, TG, HDL and CRP, and weight, Waist and BMI are measured before and after the intervention, for each patient.

General information

Acronym

IRCT registration information

IRCT registration number: **IRCT2017080135408N1**

Registration date: **2017-08-12, 1396/05/21**

Registration timing: **retrospective**

Last update:

Update count: **0**

Registration date

2017-08-12, 1396/05/21

Registrant information

Name

Masoumeh Khedri

Name of organization / entity

Ahvaz Jundishapur University of Medical Sciences

Country

Iran (Islamic Republic of)

Phone

+98 61 3292 1839

Email address

khedri.m@ajums.ac.ir

Recruitment status

Recruitment complete

Funding source

Vice chancellor for research of Ahvaz Jundishapur University of Medical Sciences

Expected recruitment start date

2016-12-05, 1395/09/15

Expected recruitment end date

2017-07-06, 1396/04/15

Actual recruitment start date

empty

Actual recruitment end date

empty

Trial completion date

empty

Scientific title

The influence of probiotics and Zinc on the serum levels of liver enzymes and lipid profile in patients with non-alcoholic fatty liver disease

Public title

The influence of probiotics and Zinc on the patients with non-alcoholic fatty liver disease

Purpose

Treatment

Inclusion/Exclusion criteria

Inclusion criteria: Patients with non-alcoholic fatty liver disease
Exclusion criteria: autoimmune hepatitis- viral hepatitis- liver metabolic diseases- liver metabolic

disease- drugs and toxins inception- age below 18- History of receiving medication for NAFLD- History of kidney or heart disease- Taking amiodarone, methotrexate, tamoxifen, glucocorticoids, sodium valproate and anti HIV drugs.

Age

From **18 years** old to **90 years** old

Gender

Both

Phase

N/A

Groups that have been masked

No information

Sample size

Target sample size: **128**

Randomization (investigator's opinion)

Randomized

Randomization description

Blinding (investigator's opinion)

Triple 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 Ahvaz Jundishpour University of Medical Sciences

Street address

Khuzestan, Ahvaz, Golestan, Ahvaz Jundishpour University of Medical Sciences

City

Ahvaz

Postal code

Approval date

2016-10-22, 1395/08/01

Ethics committee reference number

IR.AJUMS.REC.1395.435

Health conditions studied

1

Description of health condition studied

Non-alcoholic fatty liver disease

ICD-10 code

K76.0

ICD-10 code description

K76.0

Primary outcomes

1

Description

ALT

Timepoint

Before and 3 months after intervention

Method of measurement

Blood test

2

Description

AST

Timepoint

Before and 3 months after intervention

Method of measurement

Blood test

3

Description

FBS

Timepoint

Before and 3 months after intervention

Method of measurement

Blood test

4

Description

Total cholesterol

Timepoint

Before and 3 months after intervention

Method of measurement

Blood test

5

Description

Triglyceride

Timepoint

Before and 3 months after intervention

Method of measurement

Blood test

6

Description

HDL

Timepoint

Before and 3 months after intervention

Method of measurement

Blood test

7

Description

LDL

Timepoint

Before and 3 months after intervention

Method of measurement

Blood test

8

Description

CRP

Timepoint

Before and 3 months after intervention

Method of measurement

Blood test

Secondary outcomes

1

Description

Weight

Timepoint

Before and 3 months after intervention

Method of measurement

Scale

2

Description

Body mass index (BMI)

Timepoint

Before and 3 months after intervention

Method of measurement

Body mass is expressed in units of kg/m²

3

Description

Waist

Timepoint

Before and 3 months after intervention

Method of measurement

Centimeter

Intervention groups

1

Description

Two protexin probiotic capsules 600 mg (Probiotics international, United Kingdom) and two zinc sulfate capsules 25 mg (Alhavi, Iran) are used daily, at noon and at night after meals, for three months. The protexin capsule contains 7 species of probiotics. Include Lactobacillus casei, Lactobacillus rhamnosus, Streptococcus thermophilus, Lactobacillus acidophilus, Bifidobacterium breve, Bifidobacterium longum and Lactobacillus bulgaricus.

Category

Treatment - Drugs

2

Description

Two protexin probiotic capsules 600 mg (Probiotics international, United Kingdom) and two zinc placebo

capsules are used daily, at noon and at night after meals, for three months. The protexin capsule contains 7 species of probiotics. Include Lactobacillus casei, Lactobacillus rhamnosus, Streptococcus thermophilus, Lactobacillus acidophilus, Bifidobacterium breve, Bifidobacterium longum and Lactobacillus bulgaricus.

Category

Treatment - Other

3

Description

Two capsules of probiotic placebo and two zinc sulfate capsules 25 mg (Alhavi, Iran) are used daily, at noon and at night after meals, for three months.

Category

Treatment - Other

4

Description

Two probiotic placebo capsules and two zinc placebo capsules are used daily, at noon and at night after meals, for three months.

Category

Placebo

Recruitment centers

1

Recruitment center

Name of recruitment center

Imam khomeini Hospital

Full name of responsible person

Dr Masoumeh Khedri

Street address

City

Ahvaz

Sponsors / Funding sources

1

Sponsor

Name of organization / entity

Research Center for Infectious Diseases of the Digestive System, Vice Chancellor research of Ahvaz J

Full name of responsible person

Dr Behzad Sharif Makhmalzadeh

Street address

Ahvaz Jundishapur University of Medical Sciences

City

Ahvaz

Grant name

Grant code / Reference number

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

Yes

Title of funding source

Research Center for Infectious Diseases of the Digestive System, Vice Chancellor research of Ahvaz J

Proportion provided by this source

100

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****Name of organization / entity**

Ahvaz Jundishapur University of Medical Sciences

Full name of responsible person

Dr Masoumeh Khedri

Position

Hepatology and gastroenterology fellowship

Other areas of specialty/work**Street address**

Ahvaz Jundishapur University of Medical Sciences

City

Ahvaz

Postal code**Phone**

+98 61 3221 6104

Fax**Email**

khedri.m@ajums.ac.ir

Web page address

Ahvaz

Postal code**Phone**

+98 61 3221 6104

Fax

+98 61 3222 5763

Email

khedri.m@ajums.ac.ir - khedrimasoumeh@gmail.com

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

Ahvaz Jundishapur University of Medical Sciences

Full name of responsible person

Dr Masoumeh Khedri

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Full name of responsible person

Dr Masoumeh Khedri

Position

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Other areas of specialty/work**Street address**

Ahvaz Jundishapur University of Medical Sciences

City**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*