

Clinical Trial Protocol

Iranian Registry of Clinical Trials

10 Jun 2026

The effect of probiotic yogurt on blood glucose and lipid profiles serum of type 2 diabetes.

Protocol summary

Summary

Objectives: This study was designed to determine impact of Probiotic yogurt on blood glucose and lipid profiles. Inclusion criteria: Type 2 diabetes; 30-65 years; both sexes, who taking Metformin and Glibenclamide, with no renal diseases (Cr <0.4 mg/dl), high blood pressure (BP <200/110 mmHg), liver diseases. Exclusion criteria: No insulin injection; any changes in using medication; smoking; lactose intolerance; thyroid dysfunction; chronic inflammatory diseases; cardio-vascular disease; pregnancy; breast feeding; and having any weight loose or weight gain regimes. Target population were diabetic type 2 patients, sixty patients randomly divided to two intervention and control groups and respectively received 100g of probiotic yogurt (containing Lactobacillus Acidophilus and Bifidobacterium Lactis) and 100g conventional yogurt twice a day for 8 weeks in 2006.

General information

Acronym

IRCT registration information

IRCT registration number: **IRCT2016030726954N1**
Registration date: **2016-03-30, 1395/01/11**
Registration timing: **retrospective**

Last update:

Update count: **0**

Registration date

2016-03-30, 1395/01/11

Registrant information

Name

Zahra Tazakori

Name of organization / entity

Ardabil University of Medical Sciences

Country

Iran (Islamic Republic of)

Phone

+98 45 3372 8005

Email address

z.tazakori@arums.ac.ir

Recruitment status

Recruitment complete

Funding source

Proposal of Ardabil University of Medical Sciences

Expected recruitment start date

2006-03-21, 1385/01/01

Expected recruitment end date

2007-02-20, 1385/12/01

Actual recruitment start date

empty

Actual recruitment end date

empty

Trial completion date

empty

Scientific title

The effect of probiotic yogurt on blood glucose and lipid profiles serum of type 2 diabetes.

Public title

Probiotic yogurt effect on diabetes control

Purpose

Other

Inclusion/Exclusion criteria

Inclusion criteria: Type 2 diabetes; 30-65 years; both sexes, who taking Metformin and Glibenclamide, with no renal diseases (Cr <0.4 mg/dl), high blood pressure (BP <200/110 mmHg), liver diseases. Exclusion criteria: No insulin injection; any changes in using medication; smoking; lactose intolerance; thyroid dysfunction; chronic inflammatory diseases; cardio-vascular disease; pregnancy; breast feeding; and having any weight loose or weight gain regimes.

Age

From **30 years** old to **65 years** old

Gender

Both

Phase

N/A

Groups that have been masked*No information***Sample size**Target sample size: **60****Randomization (investigator's opinion)**

Randomized

Randomization description**Blinding (investigator's opinion)**

Double blinded

Blinding description**Placebo**

Used

Assignment

Parallel

Other design features**Secondary Ids**

empty

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

Ardabil University of Medical Sciences

Street addressJanbazan Square, across from Lake Shorabil, Ardabil
University of Medical Sciences**City**

Ardabil

Postal code

5618953141

Approval date

2016-03-15, 1394/12/25

Ethics committee reference number

IR.ARUMS.REC.1394.143

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

Diabetes

ICD-10 code

E11

ICD-10 code description

Non-insulin-dependent diabetes mellitus

Primary outcomes**1****Description**blood glucose (FBS) , blood glucose 2 hours after
glucose consumption**Timepoint**

before and after study

Method of measurement

photometry method pars amazon kit

2**Description**

LDL

Timepoint

before and after study

Method of measurement

Fridval formula

3**Description**

HDL,TG, total cholesterol

Timepoint

before and after study

Method of measurement

by photometry method pars amazon kit

Secondary outcomes**1****Description**

24 hours recall

Timepoint

before and after

Method of measurement

24 hours recall intake were checked by interview

Intervention groups**1****Description**Interventionl group received probiotic (100gr) twice a
day for 8 weeks**Category**

Prevention

2**Description**control group received conventional yogurt (100gr) twice
a day for 8 weeks**Category**

Prevention

Recruitment centers**1****Recruitment center****Name of recruitment center**Imam Khomeini diabetes clinic of Ardabil University of
Medical Sciences**Full name of responsible person**

Zahra Tazakori

Street address

Iran

City

Ardabil

Sponsors / Funding sources**1****Sponsor****Name of organization / entity**

Ardabil University of Medical Sciences

Full name of responsible person

Shahram Habibzadeh

Street address

Iran

City

Ardabil

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

Yes

Title of funding source

Ardabil University of Medical Sciences

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**

Ardabil University of Medical Sciences

Full name of responsible person

Zahra Tazakori

Position

PhD of Nursing and faculty member

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

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Ardabil University of Medical Sciences

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Fax**Email****Web page address****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*