

Clinical Trial Protocol

Iranian Registry of Clinical Trials

10 Jun 2026

The Effects Of Milk and Kefir Drink Consumption On Anthropometric Indises ,Lipid Profile and Blood Pressure In Premenoposal or Obese women

Protocol summary

Summary

This Randomized Controlled study is accomplished to control weight , improve lipid profile and reduce blood pressure in 75 premenoposal overweight or obese women by milk and kefir drink replacement in diet for 2 months and it's effect on anthropometric Indises , lipid profile and Blood Pressure. Particioants will be randomly devided into milk , kefir drink and control groups. Energy requirement for each person will be stimated by EER formula and the macronutrients percentage are equal for 3 groups.The milk and kefir groups consume 2 more servings of dairy products than the control group.The factors which are named above, will be measured at the beginning and at the end of the sudy.

General information

Acronym

IRCT registration information

IRCT registration number: **IRCT2013061313661N1**

Registration date: **2014-03-09, 1392/12/18**

Registration timing: **registered_while_recruiting**

Last update:

Update count: **0**

Registration date

2014-03-09, 1392/12/18

Registrant information

Name

Yasamin Fathi

Name of organization / entity

Shiraz University Of Medical Science

Country

Iran (Islamic Republic of)

Phone

+98 71 1843 5616

Email address

fathiy@sums.ac.ir

Recruitment status

Recruitment complete

Funding source

Shiraz University Of Medical sciences

Expected recruitment start date

2013-12-22, 1392/10/01

Expected recruitment end date

2014-04-22, 1393/02/02

Actual recruitment start date

empty

Actual recruitment end date

empty

Trial completion date

empty

Scientific title

The Effects Of Milk and Kefir Drink Consumption On Anthropometric Indises ,Lipid Profile and Blood Pressure In Premenoposal or Obese women

Public title

Comparision Of The Effects Of Milk and Kefir Drink Consumption On Anthropometric Indises ,Lipid Profile and Blood Pressure In Premenoposal or Obese women

Purpose

Treatment

Inclusion/Exclusion criteria

Inclusion Criteria: female sex ; age over 25 ; BMI over 25 ; agreed with lactose and mlilk ; not be menopause ; not be pregnant, not lactating, not using any drugs and nutritional supplement ; not participate in other trials over the last 6 months ; not having any disease such as thyroid ,diabetes. hypertension Exclusion Criteria : Don"t want to fallow the trial

Age

From **25 years** old to **45 years** old

Gender

Female

Phase

N/A

Groups that have been masked*No information***Sample size**Target sample size: **75****Randomization (investigator's opinion)**

Randomized

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

Not blinded

Blinding description**Placebo**

Not used

Assignment

Parallel

Other design features

Shiraz Medical University

Secondary Ids

empty

Ethics committees1**Ethics committee****Name of ethics committee**

Ethics Committee Of Shiraz Medical University

Street address

Olumpezeshki Department, Zand Street, Shiraz, Fars

City

Shiraz

Postal code**Approval date**

2013-11-29, 1392/09/08

Ethics committee reference number

1392-6751

Health conditions studied1**Description of health condition studied**

Overweight or Obese

ICD-10 code

E66

ICD-10 code description

Obesity

Primary outcomes1**Description**

Weight

Timepoint

At the beginning and at the end of the study

Method of measurement

Bascul-e- Sahand Set(with 100 grams accuracy)

2**Description**

Lipid profile

Timepoint

At the beginning and at the end of the story

Method of measurement

By use of bt 1500 set and pars test kit

3**Description**

Waist Circumference

Timepoint

At the beginning and at the end of the study

Method of measurement

Tape measure

4**Description**

Blood pressure

Timepoint

At the beginnig and at the end f the study

Method of measurement

mmgh-piezometer

Secondary outcomes1**Description**

BMI

Timepoint

At the beginning and at the end of the study

Method of measurement

weight (kg) divided by square of height (m)

Intervention groups1**Description**

Estimation of daily energy for each person by EER
 Formula Daily consumption of 500 cc milk for 2 months
 which it's energy and macronutrients has been
 calculated

Category

Lifestyle

2**Description**

Estimation Of Daily Energy for each person by EER
 Formula Daily Consumption of 750 cc kefir for 2 months
 which it's energy and macronutrients has been
 calculated.

Category

Lifestyle

3

Description

Estimation of Daily Enrgy by EER Formula Go on a diet contained 2 servings of dairy products(2servings less than intervention groups)

Category

Lifestyle

Recruitment centers

1

Recruitment center

Name of recruitment center

Cardiovascular Research Center

Full name of responsible person

Yasamin Fathi

Street address

City

Shiraz

Sponsors / Funding sources

1

Sponsor

Name of organization / entity

Shiraz Medical University

Full name of responsible person

Dr Gholamreza Hatam

Street address

Shiraz, Zand Street, Olumpezeshki Department, The 7th Floor

City

Shiraz

Grant name

Grant code / Reference number

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

Yes

Title of funding source

Shiraz Medical University

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

Shiraz Medical University, Nutrition College

Full name of responsible person

Yasamin Fathi

Position

Msc of Nutrition

Other areas of specialty/work

Street address

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Person responsible for scientific inquiries

Contact

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Full name of responsible person

Dr Shiva Faghieh

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Name of organization / entity

Medical University, Shiraz, Fars

Full name of responsible person

Yasamin Fathi

Position

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Other areas of specialty/work

Street address

City

Shiraz

Postal code

Phone

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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