

Clinical Trial Protocol

Iranian Registry of Clinical Trials

30 Jun 2026

Effect of six weeks of Pilates training and dill supplementation on serum levels of Nesfatin-1, insulin resistance with an overweight and obese woman

Protocol summary

Study aim

Comparison of the effects of Pilates exercises and dill supplementation on serum levels of Nesfatin-1 and insulin resistance and in overweight and obese women

Design

A randomized controlled clinical trial with factorial groups, triple blinded

Settings and conduct

Blood sampling of subjects in the luteal phase of menstruation after 12 hours of night fasting, 48 hours before the start of exercise program and 48 hours after the last exercise session (to prevent acute inflammation caused by exercise on biochemical variables) between 8 and 10 am Antibody in the amount of 10 ml was performed in Imam Reza Hospital (Birjand) by laboratory scientists and was stored in special tubes containing anticoagulant sodium citrate and the tube caps were closed with para-film. Blood samples for plasma separation were centrifuged and stored at minus 80 ° C for 5 minutes at 3000 rpm. Blood samples were collected for serum levels of Nesfatin-1, insulin and glucose in the pre- and post-test stages. Anthropometric characteristics were measured by the researcher at the beginning of the design and at the end of the 6 week

Participants/Inclusion and exclusion criteria

Inclusion criteria: having general physical and mental health, not participating in sports during the last 6 months, as well as diet and the use of certain herbal and chemical drugs. Exclusion criteria were cardiovascular disease, diabetes, hormonal disorders, kidney disease, liver and surgery, smoking, and any therapeutic intervention affecting the laboratory results.

Intervention groups

Pilates + dill supplement Pilates + placebo They performed Pilates exercises for 45 minutes for 6 weeks and three sessions a week. People in the dill supplement group were asked to take one dill tablet (1.1 g) after

each meal (breakfast, lunch, dinner) The placebo group received similar amounts of starch

Main outcome variables

Serum levels of Nesfatin-1 glucose

General information

Reason for update

Acronym

IRCT registration information

IRCT registration number: **IRCT20220107053655N1**

Registration date: **2022-02-05, 1400/11/16**

Registration timing: **retrospective**

Last update: **2022-02-05, 1400/11/16**

Update count: **0**

Registration date

2022-02-05, 1400/11/16

Registrant information

Name

Fatemeh Sabzevari

Name of organization / entity

The University of Birjand

Country

Iran (Islamic Republic of)

Phone

+98 56 3241 0949

Email address

fatima.sabzevari@gmail.com

Recruitment status

Recruitment complete

Funding source

Expected recruitment start date

2018-07-07, 1397/04/16

Expected recruitment end date

2018-09-07, 1397/06/16

Actual recruitment start date

empty

Actual recruitment end date

empty

Trial completion date

empty

Scientific title

Effect of six weeks of Pilates training and dill supplementation on serum levels of Nesfatin-1, insulin resistance with an overweight and obese woman

Public title

Effect of Pilates with dill supplement on obesity

Purpose

Supportive

Inclusion/Exclusion criteria**Inclusion criteria:**

Overweight and obese women (BMI more than 25 kg per square meter) with an age range of 25 to 45 years

Exclusion criteria:

Overweight and obese women (BMI more than 25 kg per square meter) with an age range of 25 to 45 years

Age

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

Gender

Female

Phase

3

Groups that have been masked

- Participant
- Care provider
- Investigator
- Outcome assessor
- Data analyser
- Data and Safety Monitoring Board

Sample size

Target sample size: **46**

Randomization (investigator's opinion)

Randomized

Randomization description

Participants in this study were Simple Random Sampling by throwing the dice

Blinding (investigator's opinion)

Triple blinded

Blinding description

Participants, data collectors, outcome assessors are blinded in this study, after informing all participants about their grouping and study objectives and they will give informed consent, they will be randomly categorized into each group. For placebo groups, capsules of the same size and the same weight as the dill capsules will be prepared. The data collectors included a laboratory expert, a blood sampler who did not know the procedure of grouping. And all participants will be treated equally on the sampling day. The statistical expert will be unaware of the participant's identities and the file containing the data and groups will be given to him/her

Placebo

Used

Assignment

Parallel

Other design features**Secondary Ids**

empty

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

Ethics committee of Birjand University

Street address

Sajjad street

City

Birjand

Province

South Khorasan

Postal code

9718993444

Approval date

2018-05-08, 1397/02/18

Ethics committee reference number

IR.BUMS.REC.1397.180

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

obesity

ICD-10 code**ICD-10 code description****Primary outcomes****1****Description**

Nesfatin-1

Timepoint

12 hours before the start of the intervention and 12 hours after the end of the intervention. Duration of intervention 6 weeks

Method of measurement

Blood test

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

Insulin resistance

Timepoint

12 hours before the start of the intervention and 12 hours after the end of the intervention. Duration of intervention 6 weeks

Method of measurement

Blood test

Intervention groups

1

Description

Intervention group: Pilates+Dill supplement. Do Pilates 3 times a week. 45 minutes per session. Take one dill (1.1g) tablet daily after each meal (breakfast, lunch, dinner)

Category

Lifestyle

2

Description

Intervention group: : Pilates+placebo. Do Pilates 3 times a week. 45 minutes per session. Take one placebo tablet daily after each meal (breakfast, lunch, dinner)

Category

Lifestyle

3

Description

Control group: : dill supplement. Take one tablet (1.1 g) daily after each meal (breakfast, lunch, dinner)

Category

Lifestyle

4

Description

Control group: placebo. Take one placebo tablet daily after each meal (breakfast, lunch, dinner)

Category

Lifestyle

Recruitment centers

1

Recruitment center

Name of recruitment center

The University of Birjand

Full name of responsible person

Fateme Sabzevari

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Sponsors / Funding sources

1

Sponsor

Name of organization / entity

The University of Birjand

Full name of responsible person

Mehdi Mogharnasi

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9717434765

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Email

public-r@bums.ac.ir

Grant name

Grant code / Reference number

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

Yes

Title of funding source

The University of Birjand

Proportion provided by this source

50

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

The University of Birjand

Full name of responsible person

Fateme Sabzevari

Position

Master graduate

Latest degree

Master

Other areas of specialty/work

Exercise Physiology

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Sharing plan

Deidentified Individual Participant Data Set (IPD)

Yes - There is a plan to make this available

Study Protocol

Yes - There is a plan to make this available

Statistical Analysis Plan

Yes - There is a plan to make this available

Informed Consent Form

Yes - There is a plan to make this available

Clinical Study Report

Yes - There is a plan to make this available

Analytic Code

Yes - There is a plan to make this available

Data Dictionary

Yes - There is a plan to make this available

Title and more details about the data/document

Part of the data such as information about the primary outcome

When the data will become available and for how long

Start of access period 6 months after printing results

To whom data/document is available

It will be available to researchers working in academic and scientific institutions

Under which criteria data/document could be used

All researchers working in in academic and scientific institutions

From where data/document is obtainable

fatima.sabzevari@gmail.com

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

After reading their request as soon as possible if the researcher find out the request valid

Comments