

Clinical Trial Protocol

Iranian Registry of Clinical Trials

02 Jul 2026

Effects of linear and daily undulating periodized resistance training on serum myokines levels, fluid balance, body composition and functional capacity in untrained women

Protocol summary

Study aim

The aim of this study was to compare the effects of linear (LP) and nonlinear (NLP) RT on serum myokines levels, fluid balance and functional capacity in overweight/obese women.

Design

The trained groups performed 3 weekly sessions for 12 weeks of resistance training with different of periodization. While the control group will continue their inactive lifestyle.

Settings and conduct

This is a blind study conducted at the University of Medical Sciences.

Participants/Inclusion and exclusion criteria

The subjects in the present study are non-athletic women with overweight / obesity, age range of 28-46, non-postmenopausal, non-disease, non-smoker, without a specific diet.

Intervention groups

In this study, we have two training groups and one control group.

Main outcome variables

Serum levels of interleukin-7, interleukin-15, insulin-like growth factor, insulin

General information

Reason for update

Acronym

IRCT registration information

IRCT registration number: **IRCT20121204011670N3**

Registration date: **2018-03-12, 1396/12/21**

Registration timing: **retrospective**

Last update: **2018-03-12, 1396/12/21**

Update count: **0**

Registration date

2018-03-12, 1396/12/21

Registrant information

Name

Mahmoud Nikseresht

Name of organization / entity

Islamic Azad University of Ilam

Country

Iran (Islamic Republic of)

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Recruitment status

Recruitment complete

Funding source

Expected recruitment start date

2017-05-22, 1396/03/01

Expected recruitment end date

2017-09-23, 1396/07/01

Actual recruitment start date

2017-07-23, 1396/05/01

Actual recruitment end date

2017-11-22, 1396/09/01

Trial completion date

empty

Scientific title

Effects of linear and daily undulating periodized resistance training on serum myokines levels, fluid balance, body composition and functional capacity in untrained women

Public title

myokines and resistance training

Purpose

Other

Inclusion/Exclusion criteria

Inclusion criteria:

Untrained women maximal oxygen uptake (VO₂max, 31–42 ml/kg/min) age (28–46 years) body mass index (26–32 kg/m²)

Exclusion criteria:

trained subjects having any kind of disease the consumption of alcohol the consumption of cigarettes having eating unusual

Age

From **28 years** old to **46 years** old

Gender

Female

Phase

3

Groups that have been masked

- Participant
- Outcome assessor
- Data analyser

Sample size

Target sample size: **45**

Actual sample size reached: **30**

Randomization (investigator's opinion)

Randomized

Randomization description

Simple random method

Blinding (investigator's opinion)

Single blinded

Blinding description

Training groups in the study are two different methods without the knowledge of other groups

Placebo

Not used

Assignment

Parallel

Other design features**Secondary Ids**

empty

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

Ethics committee of Ilam University of Medical Sciences

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Approval date

2018-02-13, 1396/11/24

Ethics committee reference number

ir.medilam, 1396.127

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

untrained women

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

Serum levels of interleukin-7

Timepoint

before and after 12 weeks of training

Method of measurement

enzyme-linked immunosorbent assay

2**Description**

Serum levels of Interleukin-15

Timepoint

before and after 12 weeks of training

Method of measurement

enzyme linked immunosorbent assay (ELISA)

3**Description**

insulin like growth factor-1

Timepoint

before and after 12 weeks of training

Method of measurement

enzyme linked immunosorbent assay (ELISA)

4**Description**

body composition

Timepoint

before and after 12 weeks of training

Method of measurement

Bio-impedance

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

fluid balance

Timepoint

before and after 12 weeks of training

Method of measurement

Bio-Impedance method

2

Description

aerobic capacity

Timepoint

before and after 12 weeks of training

Method of measurement

One-step test on the treadmill

Intervention groups

1

Description

Intervention group: Non-linear resistance training (12 weeks, three times a week for about an hour each session)

Category

Rehabilitation

2

Description

Intervention group: linear resistance training (12 weeks, three times a week for about an hour each session)

Category

Rehabilitation

3

Description

Control group: This group will maintain their sedentary lifestyle during the study period

Category

N/A

Recruitment centers

1

Recruitment center

Name of recruitment center

Islamic Azad University of Ilam Branch

Full name of responsible person

Nikseresht Mahmoud

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Sponsors / Funding sources

1

Sponsor

Name of organization / entity

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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

Ilam University of Medical Sciences

Proportion provided by this source

100

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

Islamic Azad University

Full name of responsible person

Nikseresht Mahmoud

Position

Assistant Prof.

Latest degree

Ph.D.

Other areas of specialty/work

Physiology

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

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

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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

Not applicable

Data Dictionary

Not applicable

Title and more details about the data/document

The results of this study will be published after statistical analysis

When the data will become available and for how long

One year after data collection

To whom data/document is available

Researchers working in academic and scientific institutions

Under which criteria data/document could be used

There are no special conditions

From where data/document is obtainable

Corresponding Author

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

About one month after request

Comments