

Clinical Trial Protocol

Iranian Registry of Clinical Trials

16 Sep 2026

Comparison of the effect of whole body vibration before and after eccentric exercise on functional markers of delayed onset muscle soreness

Protocol summary

Study aim

Investigating the effect of whole body vibration stimulation time on functional indicators of delayed muscle pain

Design

A controlled, parallel-group, single-blind, randomized, phase 2 clinical trial on 48 male and female seated volleyball athletes was used for randomization based on the coding method.

Settings and conduct

Subjects will be invited to the Neuromuscular Research Center of Semnan University of Medical Sciences through a researcher-made questionnaire and will be randomly assigned to one of the three intervention and control groups. It is placed on the vibration device and all measurements will be done before, immediately after, 24, 48 and 72 hours and one week and two weeks after the implementation of the protocol.

Participants/Inclusion and exclusion criteria

With 48 male and female sitting volleyball athletes, the entry and exit criteria include not consuming nutritional intervention substances (consuming high-fat food during the implementation of the protocol), drugs (not using painkillers and intramuscular injections) before the study.

Intervention groups

The first intervention group of 16 people (sitting volleyball athletes receiving whole body vibration before training), The second intervention group of 16 people (sitting volleyball athletes receiving whole body vibration after training), Control group of 16 people

Main outcome variables

Delayed contusion is measured through a visual perception chart that consists of 10 points, where 0 indicates no pain and 10 indicates pain that prevents movement. The isometric strength of the biceps muscle by the dynamometer by performing two maximum

pressures on the dynamometer while the hand is hanging by the side of the thigh. Circumference of the non-superior arm by a tape measure at the midpoint of the acromion-radial line Elbow flexion range of motion by goniometer and elbow flexion angle measurement

General information

Reason for update

Acronym

IRCT registration information

IRCT registration number: **IRCT20220713055455N1**

Registration date: **2022-11-03, 1401/08/12**

Registration timing: **registered_while_recruiting**

Last update: **2022-11-03, 1401/08/12**

Update count: **0**

Registration date

2022-11-03, 1401/08/12

Registrant information

Name

Nasrin Irvani

Name of organization / entity

Country

Iran (Islamic Republic of)

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Recruitment status

Recruitment complete

Funding source

Expected recruitment start date

2022-10-02, 1401/07/10

Expected recruitment end date

2022-11-22, 1401/09/01

Actual recruitment start date

empty

Actual recruitment end date

empty

Trial completion date

empty

Scientific title

Comparison of the effect of whole body vibration before and after eccentric exercise on functional markers of delayed onset muscle soreness

Public title

Comparison of the effect of whole body vibration before and after eccentric exercise on functional markers of delayed onset muscle soreness

Purpose

Treatment

Inclusion/Exclusion criteria**Inclusion criteria:**

Male and female seated volleyball athletes

Exclusion criteria:

Using nutritional interventions (consuming high-fat food during the implementation of the protocol) Use of interfering medicinal substances (use of painkillers and intramuscular injections) before the study

AgeFrom **20 years** old to **25 years** old**Gender**

Both

Phase

N/A

Groups that have been masked

- Participant

Sample sizeTarget sample size: **48****Randomization (investigator's opinion)**

Randomized

Randomization description

In this research, available sampling method and random grouping based on coding method will be used. The samples are coded with two-digit numbers using a random number table and randomly selected from a point of the table until the number of samples is 48.

Blinding (investigator's opinion)

Single blinded

Blinding description

In this research, 48 male and female sitting volleyball athletes are studied. The initial sampling will be voluntary, but in the next stage, subjects will be randomly (systematically) selected from among those who are eligible and registered.

Placebo

Not used

Assignment

Parallel

Other design features

-

Secondary Ids

empty

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

Ethics Committee of Semnan University of Medical Sciences

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KM 5 of Damghan road

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Semnan

Province

Semnan

Postal code

35147-99442

Approval date

2022-07-06, 1401/04/15

Ethics committee reference number

IR.SEMUMS.REC.1401.086

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

delayed onset muscle soreness

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

isometric strength

Timepoint

All measurements will be done before, immediately after, 24, 48 and 72 hours and one week and two weeks after the implementation of the protocol.

Method of measurement

To measure isometric (static) strength, the examinee holds the dynamometer in one hand so that the hand is in line with the arm and hangs next to the thigh. Without moving his hand, he displays his maximum power by pressing the dynamometer. This movement should be repeated 2 times.

2**Description**

Range of motion of the non-dominant elbow

Timepoint

All measurements will be done before, immediately after, 24, 48 and 72 hours and one week and two weeks after the implementation of the protocol.

Method of measurement

The goniometer will be used to measure the range of

motion. To measure the range of motion, the fixed arm of the goniometer will be placed parallel to the acromion-radial line and the movable arm of the goniometer will be placed on the radial-ulnar line in anatomical position.

3

Description

circumference of the non-dominant arm

Timepoint

All measurements will be done before, immediately after, 24, 48 and 72 hours and one week and two weeks after the implementation of the protocol.

Method of measurement

A tape measure is used to measure the circumference of the arm. To measure the circumference of the arm, first, the middle point of the acromion-radial line is determined, then the person places his hand in a hanging position next to the body, and the circumference of the arm is at the mentioned point. It will be measured perpendicular to the length of the arm

4

Description

muscle contusion(visual analogue scale)

Timepoint

All measurements will be done before, immediately after, 24, 48 and 72 hours and one week and two weeks after the implementation of the protocol.

Method of measurement

In order to measure muscle contusion, a visual perception chart will be used. This chart contains 10 grades, zero indicates the absence of pain, and grade 5 indicates pain that is caused by movement and causes little restriction. Grade 10 also indicates pain that may prevent any movement.

Secondary outcomes

empty

Intervention groups

1

Description

Intervention group 1: Seated volleyball athletes (16 people) after warming up and stretching for 5 minutes will perform 50 outward contractions with an intensity of 85% of 1RM and then undergo vibration therapy for one minute with a frequency of 40 Hz - with peak amplitude. They will be placed up to the top of 2 mm.

Category

Prevention

2

Description

Intervention group 2: Seated volleyball athletes (16 people) before warming up and stretching for 5 minutes will perform 50 outward contractions with an intensity of 85% of 1RM and then undergo vibration therapy for one

minute with a frequency of 40 Hz - with peak amplitude. They will be placed up to the top of 2 mm.

Category

Prevention

3

Description

Control group: The control group (16 people) - sitting volleyball athletes (16 people) will perform warm-up and stretching exercises for 5 minutes, 50 outward contractions with an intensity of 85% of 1RM, and undergo vibration therapy before and after training. are not placed

Category

Prevention

Recruitment centers

1

Recruitment center

Name of recruitment center

Neuromuscular Research Center, Faculty of Rehabilitation, Semnan University of Medical Sciences

Full name of responsible person

Fatemeh Ehsani

Street address

Tabatabai Rehabilitation Clinic, Quds Street, Mashahir Square, Semnan

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Sponsors / Funding sources

1

Sponsor

Name of organization / entity

Semnan University of Medical Sciences

Full name of responsible person

Dr. Mir Mohammad Khani

Street address

Faculty of Rehabilitation, Semnan University of Medical Sciences, 5 km Damghan Road, Semnan

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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

Semnan 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

Semnan University of Medical Sciences

Full name of responsible person

Nasrin Irvani

Position

Senior student of sports physiotherapy

Latest degree

Bachelor

Other areas of specialty/work

Physiotherapy

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Position

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Latest degree

Bachelor

Other areas of specialty/work

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Sharing plan

Deidentified Individual Participant Data Set (IPD)

Undecided - It is not yet known if there will be a plan to make this available

Study Protocol

Undecided - It is not yet known if there will be a plan to make this available

Statistical Analysis Plan

Undecided - It is not yet known if there will be a plan to make this available

Informed Consent Form

Undecided - It is not yet known if there will be a plan to make this available

Clinical Study Report

Undecided - It is not yet known if there will be a plan to make this available

Analytic Code

Undecided - It is not yet known if there will be a plan to make this available

Data Dictionary

Undecided - It is not yet known if there will be a plan to

make this available