

Clinical Trial Protocol

Iranian Registry of Clinical Trials

11 Sep 2026

The effect of a Falling Safely Training (FAST) protocol on head and hip acceleration during the fall, some components of gait and quality of life in older adults

Protocol summary

Study aim

The aim of the study is to investigate the effect of a falling safely train (FAST) course on head and hip acceleration during a fall, some gait components, and quality of life in the older adults.

Design

Randomized clinical trial, with experimental and control group, single-blind, with pre-test and post-test on 28 participants. A simple lottery method is used for randomization

Settings and conduct

The tests will be performed in a 3D motion analysis laboratory and quality of life will be assessed with a questionnaire. The exercises will be performed in gyms. For blinding, each participant will assume that everyone is doing the same activity as him.

Participants/Inclusion and exclusion criteria

Entry conditions: elderly people 60 to 70 years old; with no history of sports; with hip bone mineral density ≥ -2.3 (i.e. t-score ≥ -2.3); walking independently. Not entering conditions: currently undergoing physiotherapy treatment and taking anticoagulants; In the last 3 years, any injury or abnormality including any fractures or musculoskeletal surgeries, heart diseases and diabetes.

Intervention groups

The falling safely train is a progressive exercise that aims to teach how to effectively reduce impact forces during falls, change the angles of the trunk and knees during landing to reduce body acceleration, and use rolling movements to optimally distribute impact forces applied to each part of the body along the path of contact with the ground. The effect of this type of training on gait components and quality of life in the elderly is also examined. The experimental group performs the above exercises for 6 weeks but the control group only performs the pre-test and post-test.

Main outcome variables

It is predicted that head and hip acceleration during falls will decrease and gait components (step length, stride width, gait speed) and quality of life of the elderly will improve.

General information

Reason for update

Acronym

IRCT registration information

IRCT registration number: **IRCT20260602069619N1**

Registration date: **2026-06-18, 1405/03/28**

Registration timing: **prospective**

Last update: **2026-06-18, 1405/03/28**

Update count: **0**

Registration date

2026-06-18, 1405/03/28

Registrant information

Name

Fatemeh Khosravani

Name of organization / entity

The University of Kharazmi

Country

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Recruitment status

Recruitment complete

Funding source

Expected recruitment start date

2026-07-06, 1405/04/15

Expected recruitment end date

2026-08-23, 1405/06/01

Actual recruitment start date

empty

Actual recruitment end date

empty

Trial completion date

empty

Scientific title

The effect of a Falling Safely Training (FAST) protocol on head and hip acceleration during the fall, some components of gait and quality of life in older adults

Public title

Effect of falling safely train on older adult

Purpose

Education/Guidance

Inclusion/Exclusion criteria**Inclusion criteria:**

Bone mineral density of the hip ≥ -2.3 Independent walking (no use of walking aids)

Exclusion criteria:

Damage or abnormality that affecting on testing process
They are currently undergoing physiotherapy treatment

Age

From **60 years** old to **70 years** old

Gender

Both

Phase

N/A

Groups that have been masked

- Participant

Sample size

Target sample size: **28**

Randomization (investigator's opinion)

Randomized

Randomization description

Simple random sampling: Lottery method. Each participant is assigned a number and written on a piece of paper. Totally 28 papers. The papers are placed in a bag. Then we take out the papers one by one randomly, the first 14 papers are placed in the experimental group and the next 14 papers are placed in the control group.

Blinding (investigator's opinion)

Single blinded

Blinding description

The grouping of participants into two experimental and control groups is done randomly by the researcher after the pre-test. Participants do not know which group they may be placed in during the pre-test.

Placebo

Not used

Assignment

Other

Other design features**Secondary Ids**

empty

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

Ethics committee of Kharazmi university

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Next to Shahid Keshvari Sports Complex, Intersection of Kosha and Varzesh Streets, Mirdamad Blvd

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1544733111

Approval date

2026-01-07, 1404/10/17

Ethics committee reference number

IR.KHU.REC.1404.143

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

Prevention of fall injuries in the older adults

ICD-10 code

R29.6

ICD-10 code description

Repeated falls

Primary outcomes**1****Description**

Head and hip acceleration in fall test.

Timepoint

Head and hip acceleration during a fall is measured at the beginning of the study and after 6 weeks of training to assess changes.

Method of measurement

3D motion analysis system

2**Description**

Gait components include stride length, stride width, and walking speed.

Timepoint

Measurements will be taken at the beginning of the study and again after 6 weeks of training.

Method of measurement

3D motion analysis system

3**Description**

Quality of life in older adults

Timepoint

The assessment will be conducted at the beginning of

Person responsible for scientific inquiries

Contact

Name of organization / entity

The Kharazmi University

Full name of responsible person

Seyed Sadredin Shojaedin

Position

Professor

Latest degree

Ph.D.

Other areas of specialty/work

Corrective exercises

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Person responsible for updating data

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Student

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Sharing plan

Deidentified Individual Participant Data Set (IPD)

No - There is not a plan to make this available

Justification/reason for indecision/not sharing IPD

To protect participant privacy

Study Protocol

No - There is not a plan to make this available

Statistical Analysis Plan

No - There is not a plan to make this available

Informed Consent Form

No - There is not a plan to make this available

Clinical Study Report

No - There is not a plan to make this available

Analytic Code

Not applicable

Data Dictionary

No - There is not a plan to make this available