

Clinical Trial Protocol

Iranian Registry of Clinical Trials

10 Jun 2026

A randomized controlled trial to comparative study of the effect of single and dual cognitive task-oriented balance exercises on postural control and functional balance in subjects with chronic stroke

Protocol summary

Study aim

The aim of this study is to comparative study of the effect of single and dual cognitive task-oriented balance exercises on postural control, functional balance and mobility and cognitive function in subjects with chronic stroke.

Design

This study is a controlled randomized trial with parallel groups and blinded outcome assessment. Fifty-four chronic stroke survivors will participate in this study and they will be randomly assigned to three groups: control group, single task-oriented balance exercises group and cognitive dual task-oriented balance exercises group. Randomization will be performed by the person who is not involved in the study using sequentially numbered sealed envelopes.

Settings and conduct

This study will be performed in Djavad Mowafaghian Research Centre of Intelligent Neuro-Rehabilitation Technologies. Control group receives conventional rehabilitation. Single and cognitive dual task-oriented balance exercises groups receive 24 sessions of task-oriented balance exercises (8 weeks, 3 sessions per week, 60 to 90 minutes per session) in addition to conventional rehabilitation. In single task-oriented balance exercises group, the attention of patient will be focused on maintaining his/her balance while in the cognitive dual task-oriented balance exercises group, the attention of patient will be focused on maintaining his/her balance while in the cognitive dual task-oriented balance exercises group, the attention of patient will be focused concurrently on maintaining balance and performing cognitive tasks. Assessment of the primary and secondary outcomes is performed by individuals who are blinded to the groups allocation before and after receiving task-oriented balance exercises and 8 weeks after receiving these exercises in the intervention groups

and before and after conventional rehabilitation and 8 weeks after conventional rehabilitation in the control group.

Participants/Inclusion and exclusion criteria

The main inclusion criteria include having the first experience of stroke; Having the ability to perform the most difficult condition of laboratory test (quiet standing on foam surface with closed eyes while performing cognitive task); Having an acceptable level of cognitive function, i.e. score equal to or greater than 23 on the Mini Mental State Examination. Subjects are excluded in the case of having unilateral visuospatial neglect and co-morbid other neurological diseases, orthopedic disorders, diabetes or addiction.

Intervention groups

Control group; Single task-oriented balance exercise; Cognitive dual task-oriented balance exercise

Main outcome variables

Functional balance; Postural control

General information

Reason for update

Acronym

IRCT registration information

IRCT registration number: **IRCT20140304016830N9**

Registration date: **2018-02-25, 1396/12/06**

Registration timing: **registered_while_recruiting**

Last update: **2018-02-25, 1396/12/06**

Update count: **0**

Registration date

2018-02-25, 1396/12/06

Registrant information

Name

Ghorban Taghizadeh

Name of organization / entity

School of Rehabilitation Sciences, Iran University of

Medical

Country

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Recruitment status

Recruitment complete

Funding source**Expected recruitment start date**

2018-02-04, 1396/11/15

Expected recruitment end date

2018-06-21, 1397/03/31

Actual recruitment start date

empty

Actual recruitment end date

empty

Trial completion date

empty

Scientific title

A randomized controlled trial to comparative study of the effect of single and dual cognitive task-oriented balance exercises on postural control and functional balance in subjects with chronic stroke

Public title

Effect of task-oriented balance exercises on balance of patients with chronic stroke

Purpose

Treatment

Inclusion/Exclusion criteria**Inclusion criteria:**

Having the first experience of stroke Passing 6-24 months since stroke Having an ability to walk at least 10 meters without assistive devices Having the ability to perform the most difficult condition of laboratory test (quiet standing on foam surface with closed eyes while performing cognitive task) Having an acceptable level of cognitive function, i.e. score equal to or greater than 23 on the Mini Mental State Examination

Exclusion criteria:

Having unilateral visuospatial neglect (i.e., obtaining score less than 44 at star cancellation test) Co-morbid other neurological diseases Having orthopedic disorders (such as low back pain, arthritis and flat foot) Having diabetes according to the report of patient, or the patient's family or physician Having addiction according to the report of patient, or the patient's family or physician

Age

From 35 years old to 65 years old

Gender

Both

Phase

N/A

Groups that have been masked

- Outcome assessor

Sample size

Target sample size: 54

Randomization (investigator's opinion)

Randomized

Randomization description

Randomization will be performed by the person who is not involved in the study using sequentially numbered sealed envelopes.

Blinding (investigator's opinion)

Single blinded

Blinding description

Evaluation of the primary and secondary outcomes is performed by experienced individuals who are blind on the groups allocation.

Placebo

Not used

Assignment

Parallel

Other design features**Secondary Ids**

empty

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

Ethics committee of Iran University of Medical Sciences

Street address

Iran University of Medical Sciences, Shahid Hemmat Highway, Tehran

City

Tehran

Province

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Postal code

۱۴۳۹۶۱۴۵۳۵

Approval date

2017-12-08, 1396/09/17

Ethics committee reference number

IR.IUMS.REC 1396.30275

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

stroke

ICD-10 code

I64

ICD-10 code description

Stroke, not specified as haemorrhage or infarction

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

Functional balance

Timepoint

Before and after intervention and 8 weeks after intervention

Method of measurement

Berg Balance Scale Questionnaire, Tinetti Balance test

2

Description

Postural control

Timepoint

Before and after intervention and 8 weeks after intervention

Method of measurement

Force plate (postural sway parameters)

Secondary outcomes

1

Description

Functional mobility

Timepoint

Before and after intervention and 8 weeks after intervention

Method of measurement

Single and dual Timed Up & Go Test, Dynamic gait index Questionnaire

2

Description

Cognitive function

Timepoint

Before and after intervention and 8 weeks after intervention

Method of measurement

Wechsler memory scale

Intervention groups

1

Description

Control group: receives conventional rehabilitation (including exercises for upper and lower limb stretching, range of motion and activities of daily living).

Category

Rehabilitation

2

Description

First intervention group: Single task-oriented balance exercises group that receives 24 sessions of single task-oriented balance exercises (8 weeks, 3 sessions per week, 60 to 90 minutes per session) in addition to the conventional rehabilitation. Single task-oriented balance exercises include three types of standing, transferring and walking activities which performed based on two principals of difficulty level of motor task and individual

safety.

Category

Rehabilitation

3

Description

Second Intervention group: cognitive dual task-oriented balance exercises group that receives 24 sessions of cognitive dual task-oriented balance exercises (8 weeks, 3 sessions per week, 60 to 90 minutes per session) in addition to the conventional rehabilitation. In this group, task-oriented balance exercises (including three types of standing, transferring and walking activities based on two principals of difficulty level of motor task and individual safety) are performed simultaneously with cognitive tasks (e.g., backward counting, saying the days of the week and the months of the year in reverse order, etc.).

Category

Rehabilitation

Recruitment centers

1

Recruitment center

Name of recruitment center

Rehabilitation centers and hospitals in Tehran

Full name of responsible person

Ghorban Taghizadeh

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Sponsors / Funding sources

1

Sponsor

Name of organization / entity

Iran University of Medical Sciences

Full name of responsible person

Vice Chancellor for research of Iran University of Medical Sciences, Dr. Seyed Kazem Malakouti

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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

Iran 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**

Iran University of Medical Sciences

Full name of responsible person

Ghorban Taghizadeh

Position

Assistant Professor

Latest degree

Ph.D.

Other areas of specialty/work

Neuroscience

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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

After the completion of this study, a manuscript regarding the documentation and the results of the study will be prepared and published. If more details are needed, individuals can send their request by email to the person responsible for scientific accountability.

When the data will become available and for how long

8 months after publishing the results

To whom data/document is available

Researchers working in academic and scientific institutions

Under which criteria data/document could be used

Use of the documentation is permitted upon written permission.

From where data/document is obtainable

Ghorban Taghizadeh Address: Iran University of Medical Sciences, Shahid Hemmat Highway, Tehran Tel: 00982122227124 E-mail: taghizadeh.gh@iums.ac.ir

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

Just sending a request by email and mentioning the explanation about the cause of the need for documentation is enough.

Comments