

Clinical Trial Protocol

Iranian Registry of Clinical Trials

27 Jun 2026

Comparison of the Effects of Cognitive_Balance Dual-Task Training and Single Task Balance Training on Static Balance of People with Anterior Cruciate Ligament Reconstruction (ACL_R)

Protocol summary

Study aim

The main purposes of this study are the comparison of some of center of pressure (COP) parameters in static balance task while standing on reconstructed limb before and after intervention in each group and between groups, also survey of correlation between COP parameters and level of functional disability measured by KOOS questionnaire in people with anterior cruciate ligament reconstruction.

Design

Clinical trial with parallel control group, double blind and randomized grouping, 15 people in each group

Settings and conduct

This study will be carried out at the Biomechanics Lab of rehabilitation sciences faculty of Tabriz University of Medical Sciences. Individuals in intervention group will do progressive balance training simultaneously with cognitive tasks for 20 minutes 3 sessions a week for 6 weeks. The control group will just do balance training. Participants, researcher, and data recorder and analyzer will be blind to groups.

Participants/Inclusion and exclusion criteria

In this study the people with unilateral anterior cruciate ligament reconstructed by hamstring tendon allograft and ability of full weight bearing on reconstructed limb and at least 120 degree reconstructed knee range of motion will participate; and they will be eliminated if they have a history of neurological, vestibular and visual disorder, injury or surgery of the spine or other segments of involved limb or other ligaments of the reconstructed knee.

Intervention groups

Intervention group will do balance training and cognitive tasks simultaneously (Dual Task). However, control group will just do balance training (Single Task).

Main outcome variables

COP parameters include sway amplitude, velocity, path

length and Mean Square (RMS) amplitude and velocity of COP in anterior-posterior and medial-lateral direction, total path length of COP ,Root , ellipse sway area and KOOS questionnaire score.

General information

Reason for update

Acronym

Anterior Cruciate Ligament Reconstruction (ACL_R)

IRCT registration information

IRCT registration number: **IRCT20180925041138N1**

Registration date: **2019-05-31, 1398/03/10**

Registration timing: **prospective**

Last update: **2019-05-31, 1398/03/10**

Update count: **0**

Registration date

2019-05-31, 1398/03/10

Registrant information

Name

Jalal Ahadi

Name of organization / entity

Country

Iran (Islamic Republic of)

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Recruitment status

Recruitment complete

Funding source

Expected recruitment start date

2019-06-22, 1398/04/01

Expected recruitment end date

2020-03-19, 1398/12/29

Actual recruitment start date

empty

Actual recruitment end date

empty

Trial completion date

empty

Scientific title

Comparison of the Effects of Cognitive_Balance Dual-Task Training and Single Task Balance Training on Static Balance of People with Anterior Cruciate Ligament Reconstruction (ACL_R)

Public title

Affect of Dual_Task Balance Training on Balance of People with Anterior Cruciate Ligament Reconstruction

Purpose

Treatment

Inclusion/Exclusion criteria**Inclusion criteria:**

Unilateral ACL Reconstruction Surgery by Hamstring Tendon Allograft Full Weight Bearing Ability on Reconstructed limb At Least 120 Degree Reconstructed Knee Range of Motion

Exclusion criteria:

The History of Neurological, Vestibular and Visual Disorder The History of Injury or Surgery of the Spine The History of Injury or Surgery of the Other Segments of Reconstructed Limb The History of Injury or Surgery of the Other Ligaments of Reconstructed Knee

Age

From **18 years** old to **45 years** old

Gender

Both

Phase

N/A

Groups that have been masked

- Participant
- Investigator
- Outcome assessor
- Data analyser

Sample size

Target sample size: **30**

Randomization (investigator's opinion)

Randomized

Randomization description

In this study people with inclusion criteria will be divided into control and Intervention groups according to simple randomization way by Random Software Allocation.

Blinding (investigator's opinion)

Double blinded

Blinding description

In this study all of the participants will be unaware of group of study they belong to. Also researcher, outcomes assessors and data analyzer will be unaware the group each participants belong to. Inevitably, the trainer physiotherapist base on intervention type will not be blind. However, he will not be allowed to outcome assessment and data analysis.

Placebo

Not used

Assignment

Parallel

Other design features**Secondary Ids**

empty

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

Ethics committee of Tabriz University of Medical Sciences

Street address

Tabriz University of Medical Sciences, Golgasht Street, Azadi Street, Tabriz

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Postal code

5166615669

Approval date

2019-01-21, 1397/11/01

Ethics committee reference number

IR.TBZMED.REC.1397.865

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

Anterior Cruciate Ligament Reconstruction

ICD-10 code

S83.51

ICD-10 code description

Sprain of anterior cruciate ligament of knee

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

Mean Anterior - Posterior Sway amplitude of Center of Pressure

Timepoint

Before starting balance training and at the end of treatment (18 session later)

Method of measurement

Force Plate Instrument

2**Description**

Mean Medial - Lateral Sway amplitude of Center of Pressure

Timepoint

Before starting balance training and at the end of treatment (18 session later)

Method of measurement

Force Plate Instrument

3**Description**

Total path length of Center of Pressure

Timepoint

Before starting balance training and at the end of treatment (18 session later)

Method of measurement

Force Plate Instrument

4**Description**

Root-mean-square amplitude of center of pressure in anterior - posterior direction

Timepoint

Before starting balance training and at the end of treatment (18 session later)

Method of measurement

Force Plate Instrument

5**Description**

Root-mean-square amplitude of center of pressure in medial - lateral direction

Timepoint

Before starting balance training and at the end of treatment (18 session later)

Method of measurement

Force Plate Instrument

6**Description**

Root-mean-square velocity of center of pressure in anterior - posterior direction

Timepoint

Before starting balance training and at the end of treatment (18 session later)

Method of measurement

Force Plate Instrument

7**Description**

Root-mean-square velocity of center of pressure in medial - lateral direction

Timepoint

Before starting balance training and at the end of treatment (18 session later)

Method of measurement

Force Plate Instrument

8**Description**

Ellipse sway area

Timepoint

Before starting balance training and at the end of

treatment (18 session later)

Method of measurement

Force Plate Instrument

9**Description**

Center of pressure velocity in anterior - posterior direction

Timepoint

Before starting balance training and at the end of treatment (18 session later)

Method of measurement

Force Plate Instrument

10**Description**

Center of pressure velocity in medial - lateral direction

Timepoint

Before starting balance training and at the end of treatment (18 session later)

Method of measurement

Force Plate Instrument

11**Description**

Path length of center of pressure in anterior - posterior direction

Timepoint

Before starting balance training and at the end of treatment (18 session later)

Method of measurement

Force Plate Instrument

12**Description**

Path length of center of pressure in medial - lateral direction

Timepoint

Before starting balance training and at the end of treatment (18 session later)

Method of measurement

Force Plate Instrument

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

The Knee injury and Osteoarthritis Outcome Score (KOOS) questionnaire

Timepoint

Before starting balance training and at the end of treatment (18 session later)

Method of measurement

Force Plate Instrument

Intervention groups

1

Description

Intervention group: Anterior cruciate reconstructed patients during rehabilitation in balance training phase will do progressive balance training including single leg standing on reconstructed limb, standing on bi-direction balance board in med-lat direction, standing on bi-direction balance board in ant-post direction, standing on bi-direction balance board with foam surface(10 cm thickness), standing on multi-direction balance board, standing on multi-direction balance board with foam surface(10 cm thickness) simultaneously with cognitive tasks including backward counting of numbers at a specified interval, saying the days of the week in reverse and alternative ways, saying the names started with special letters, reverse words spelling (Dual Task) for 20 minutes, 3 sessions a week for 6 weeks.

Category

Rehabilitation

2

Description

Control group: Anterior cruciate reconstructed patients during rehabilitation in balance training phase will do progressive balance training including single leg standing on reconstructed limb, standing on bi-direction balance board in med-lat direction, standing on bi-direction balance board in ant-post direction, standing on bi-direction balance board with foam surface(10 cm thickness), standing on multi-direction balance board, standing on multi-direction balance board with foam surface(10 cm thickness) without any cognitive tasks (Single Task) for 20 minutes, 3 sessions a week for 6 weeks.

Category

Rehabilitation

Recruitment centers

1

Recruitment center

Name of recruitment center

Shohada medical research & training hospital

Full name of responsible person

Dr. Amin Moradi

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Sponsors / Funding sources

1

Sponsor

Name of organization / entity

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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

Tabriz 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

Tabriz University of Medical Sciences

Full name of responsible person

Jalal Ahadi

Position

Assistance Professor

Latest degree

Ph.D.

Other areas of specialty/work

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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

There is no further information.

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

No - There is not a plan to make this available

Data Dictionary

No - There is not a plan to make this available