

Clinical Trial Protocol

Iranian Registry of Clinical Trials

04 Jul 2026

The effect of dynamic neuromuscular stabilization exercises on stress and combined urinary incontinence in women with multiple sclerosis

Protocol summary

Study aim

The effect of dynamic neuromuscular stabilization exercises on urinary incontinence in patients with multiple sclerosis

Design

In this study, due to the small number of samples, in consultation with a statistician, it was decided to use the random blocking method. Blocking is usually used to balance the number of samples assigned to each of the groups studied. The statistician used the website <https://www.sealedenvelope.com/simple-randomiser/v1/lists>, considering that we had 2 groups, to design a suitable blocking table. Divides patients according to this table is unaware of the study.

Settings and conduct

The place of this study is Isfahan, Ayatollah Kashani St., Kashani Hospital, physiotherapy ward and the time of its conduct is 2021. Study begins with obtaining permission from the ethics committee. Volunteers attend the physiotherapy department of Kashani Hospital. All stages of the work and the purpose of the research for patients are described in detail and the consent form is completed by them. In this study, patients are unaware of the groups. In both experimental and control groups, exercise therapy is performed, but the exercises are different. Also, the person doing the assessments is unaware of the groups.

Participants/Inclusion and exclusion criteria

Women diagnosed with multiple sclerosis (by a neurologist) who have urinary incontinence. Inclusion criteria include having a low disability to participate in the study and having at least one urinary incontinence in the past month. Exclusion from the study is other neurological diseases.

Intervention groups

In this study, the control group of Kegel exercises and the experimental group of dynamic neuromuscular stabilization receive.

Main outcome variables

Pelvic floor muscle strength, urinary incontinence, quality of life

General information

Reason for update

Acronym

IRCT registration information

IRCT registration number: **IRCT20200101045970N5**

Registration date: **2021-10-27, 1400/08/05**

Registration timing: **registered_while_recruiting**

Last update: **2021-10-27, 1400/08/05**

Update count: **0**

Registration date

2021-10-27, 1400/08/05

Registrant information

Name

Ehsan Ghasemi

Name of organization / entity

Country

Iran (Islamic Republic of)

Phone

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Recruitment status

Recruitment complete

Funding source

Expected recruitment start date

2021-09-23, 1400/07/01

Expected recruitment end date

2022-03-16, 1400/12/25

Actual recruitment start date

empty

Actual recruitment end date

empty

Trial completion date

empty

Scientific title

The effect of dynamic neuromuscular stabilization exercises on stress and combined urinary incontinence in women with multiple sclerosis

Public title

The effect of stabilizing exercises on urinary incontinence in patients with MS

Purpose

Treatment

Inclusion/Exclusion criteria**Inclusion criteria:**

Women between the ages of 18 and 55. Diagnosis of multiple sclerosis relapsing-remitting whose disease condition has stabilized (in the last 4 months. No change in symptoms) According to the EDSS questionnaire, their disability should be less than or equal to 5.5. Women who have experienced stress incontinence at least once in the past month.

Exclusion criteria:

Pregnancy Existence of any neurological disease (except multiple sclerosis) or kidney disease History of childbirth in the last 6 months History of gastrointestinal surgery and abdominal surgery in less than 6 months Infection of the lower urinary tract Recurrence of the disease during the project Existence of musculoskeletal pain such as back pain Taking the drug for urinary incontinence (in case of discontinuation of the drug can be included in the study with the opinion of a doctor) Being a virgin

Age

From **18 years** old to **55 years** old

Gender

Female

Phase

N/A

Groups that have been masked

- Participant
- Outcome assessor

Sample size

Target sample size: **30**

Randomization (investigator's opinion)

Randomized

Randomization description

In this study, due to the small number of samples, in consultation with a statistician, it was decided to use the random blocking method. Blocking is usually used to balance the number of samples assigned to each of the groups studied. The statistician used the website <https://www.sealedenvelope.com/simple-randomiser/v1/lists>, considering that we had 2 groups, to design a suitable blocking table. Divides patients according to this table is unaware of the study.

Blinding (investigator's opinion)

Double blinded

Blinding description

In this study, patients are unaware of the groups. In this study, exercise therapy is presented in both experimental and control groups, but they are different.

In this study, patients try to come at different times and do not even contact each other so that they do not know the details of each other's exercises. Also, the person doing the assessments is unaware of the groups.

Placebo

Not used

Assignment

Parallel

Other design features**Secondary Ids**

empty

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

Faculty of Nursing, Management and Rehabilitation - Isfahan University of Medical Sciences (Research

Street address

Hezar Jirib

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Postal code

81746-73461

Approval date

2021-10-01, 1400/07/09

Ethics committee reference number

IR.MUI.NUREMA.REC.1400.130

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

Multiple Sclerosis

ICD-10 code

G35

ICD-10 code description

Multiple sclerosis

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

Pelvic floor muscle strength

Timepoint

Before treatment, after 3 weeks from the beginning of the sessions, after the end of the sessions and 2 months after the end of the last treatment session

Method of measurement

The modified Oxford Scale, a 6-point scale, is used to assess pelvic floor muscle strength.

2

Description

The rate of urinary incontinence

Timepoint

Before treatment, after 3 weeks from the beginning of the sessions, after the end of the sessions and 2 months after the end of the last treatment session

Method of measurement

The severity index of urinary incontinence in women is used to assess the rate of urinary incontinence

Secondary outcomes

1

Description

Quality of Life

Timepoint

Before treatment, after 3 weeks from the beginning of the sessions, after the end of the sessions and 2 months after the end of the last treatment session

Method of measurement

The LUTS-QOL questionnaire is used to assess the effect of urinary incontinence on the quality of life of these people.

Intervention groups

1

Description

Intervention group: Receive dynamic neuromuscular stabilization exercises. These exercises are performed in different positions that are actually modeled on the positions of the growing baby. Exercises start with simple situations and gradually become more complex. In fact, the exercises start with the supine and then progress to the rolling, sitting, bear and squat-like postures. The workout lasts for 6 weeks. The patient goes to physiotherapy twice a week.

Category

Rehabilitation

2

Description

Control group: Receive Kegel exercises. This exercise involves the patient contracting the pelvic floor muscles under the supervision of a therapist. In each session, moderate and intense exercises are performed. Moderate-intensity exercises last between 6 and 10 seconds, and high-intensity exercises last between 1 and 3 seconds. The exercises start from the supine position and are gradually taught in different positions such as sitting and standing. The patient is treated twice a week for 6 weeks.

Category

Rehabilitation

Recruitment centers

1

Recruitment center

Name of recruitment center

Kashani Hospital

Full name of responsible person

Ehsan Ghasemi

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kashani

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Sponsors / Funding sources

1

Sponsor

Name of organization / entity

Esfahan University of Medical Sciences

Full name of responsible person

shaghayegh haghjoo

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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

Esfahan 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

Esfahan University of Medical Sciences

Full name of responsible person

Ehsan Ghasemi

Position

Assistant professor

Latest degree

Ph.D.

Other areas of specialty/work

Physiotherapy

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Person responsible for scientific inquiries

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Full name of responsible person

Ehsan Ghasemi

Position

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Latest degree

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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

Because of confidentiality

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

No - There is not a plan to make this available

Data Dictionary

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

Title and more details about the data/document

-

When the data will become available and for how long

-

To whom data/document is available

-

Under which criteria data/document could be used

-

From where data/document is obtainable

-

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

-

Comments