

# Clinical Trial Protocol

## Iranian Registry of Clinical Trials

09 Jun 2026

### Efficacy of Postural Restoration exercises on pain, ROM and disability of the patients with chronic non-specific low back pain

#### Protocol summary

##### Study aim

1. Comparing the intensity of pain between groups 2. Comparing the amount of disability between groups using Oswestry disability questionnaire 3. Comparing passive hip adduction, abduction, internal rotation and external rotation ROM between groups 4. Comparing passive shoulder internal and external rotation ROM between groups 5. Comparing Gleno-humeral internal rotation deficiency(GIRD) between groups 6. Assessment of relation between hip and contralateral shoulder ROM between groups

##### Design

Single blinded randomized controlled trial, pre-test post-test design. with block Randomization method in 18 patients with chronic non-specific low back pain

##### Settings and conduct

The process has been performed in a private clinic. Patients were allocated into control and intervention groups and randomized by block randomization method. After meeting the inclusion criteria, patients were evaluated at first session and sixth session. Patients in the treatment group were asked to perform five Postural restoration exercises and patients in the control group were asked to perform five routine exercises. The process lasted 5 sessions and patients were reevaluated in the sixth session.

##### Participants/Inclusion and exclusion criteria

1. Patients who had positive Ober`s test on one side indicating pelvic girdle malalignment, and limited contralateral shoulder internal rotation ROM. 2. Patients who didn`t have specific spine pathology like spondylolisthesis, canal stenosis, fracture and infection. 3. At least 3 months had pasted from the first episode of low back pain. 4. Patients who didn`t have any neurologic sign and symptoms like radicular pain, numbness and paresthesia

##### Intervention groups

Control group: Routine physiotherapy exercise for low back pain Intervention group: Postural restoration

exercise

##### Main outcome variables

Pain Disability(Oswestry disability questionnaire) Hip ROM Shoulder ROM

#### General information

##### Reason for update

##### Acronym

##### IRCT registration information

IRCT registration number: **IRCT20201230049883N1**

Registration date: **2021-01-18, 1399/10/29**

Registration timing: **retrospective**

Last update: **2021-01-18, 1399/10/29**

Update count: **0**

##### Registration date

2021-01-18, 1399/10/29

##### Registrant information

##### Name

Mehran Fathizadeh

##### Name of organization / entity

##### Country

Iran (Islamic Republic of)

##### Phone

+98 21 6609 8062

##### Email address

m3hran.fathi@gmail.com

##### Recruitment status

**Recruitment complete**

##### Funding source

##### Expected recruitment start date

2019-07-01, 1398/04/10

##### Expected recruitment end date

2020-05-19, 1399/02/30

##### Actual recruitment start date

2019-07-01, 1398/04/10

**Actual recruitment end date**

2020-02-20, 1398/12/01

**Trial completion date**

2020-02-20, 1398/12/01

**Scientific title**

Efficacy of Postural Restoration exercises on pain, ROM and disability of the patients with chronic non-specific low back pain

**Public title**

Efficacy of Postural Restoration exercises on patients with chronic non-specific low back pain

**Purpose**

Treatment

**Inclusion/Exclusion criteria****Inclusion criteria:**

Patients with pelvic malalignment determined by Ober`s test  
Patients with chronic low back pain who has lasted at least 3 months  
Patients who don`t have specific pathology causing low back pain like tumors, fractures, infection, spondylolisthesis, canal stenosis  
Patients who don`t have previous surgery  
BMI less than 30  
Patients who don`t have neurologic sign and symptoms like radicular pain or numbness in lower extremities  
Patients who are not drug addicted  
Patients who are not suffering from mental disorders  
Patients who are not pregnant  
Patients who don`t have neurological disease like MS or CVA  
Patients who don`t have rheumatologic disease

**Exclusion criteria:**

Inability of the patients to learn the exercises  
Bilateral positive Ober`s test  
Using pain killers

**Age**

From **20 years** old to **45 years** old

**Gender**

Both

**Phase**

N/A

**Groups that have been masked**

- Participant

**Sample size**

Target sample size: **16**

Actual sample size reached: **18**

**Randomization (investigator's opinion)**

Randomized

**Randomization description**

By the use of software, block randomization method was used. There were 5 blocks and patients equally devoted to both control(A) and intervention(B) groups at each block.

**Blinding (investigator's opinion)**

Single blinded

**Blinding description**

Patients were blind to the group(control/intervention) that they were in.

**Placebo**

Not used

**Assignment**

Parallel

**Other design features****Secondary Ids**

empty

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

Ethics committee of university of social welfare and rehabilitation sciences

**Street address**

Koodakyar Ave, Daneshjoo Bly, Evin, Tehran, Iran

**City**

Tehran

**Province**

Tehran

**Postal code**

1985713834

**Approval date**

2019-06-26, 1398/04/05

**Ethics committee reference number**

IR.USWR.REC.1398.034

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

Chronic non-specific low back pain

**ICD-10 code**

M54.5

**ICD-10 code description**

Low back pain

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

Intensity of pain

**Timepoint**

Before intervention and 12 days later

**Method of measurement**

Visual analog scale

**2****Description**

Hip internal and external ROM

**Timepoint**

Before intervention and 12 days later

**Method of measurement**

Standard goniometer

**3****Description**

Hip abduction and adduction ROM

**Timepoint**

Before intervention and 12 days later

## Method of measurement

Inclinometer

### 4

#### Description

Score of disability in Oswestry disability questionnaire

#### Timepoint

Before intervention and 12 days later

#### Method of measurement

Oswestry disability questionnaire

## Secondary outcomes

### 1

#### Description

Shoulder internal and external rotation ROM

#### Timepoint

Before intervention and 12 days later

#### Method of measurement

Standard Goniometer

## Intervention groups

### 1

#### Description

Intervention group: Postural restoration exercises

#### Category

Treatment - Other

### 2

#### Description

Control group: Routine physiotherapy exercises for low back pain

#### Category

Treatment - Other

## Recruitment centers

### 1

#### Recruitment center

##### Name of recruitment center

Private Clinic (Dayani\_physio\_clinic)

##### Full name of responsible person

Mohammad Reza Dayani

##### Street address

Unit 8, No 193, Mollasadra street, Tehran, Iran

##### City

Tehran

##### Province

Tehran

##### Postal code

0000000000

##### Phone

+98 21 8861 3954

##### Email

m3hran.fathi@gmail.com

## Sponsors / Funding sources

### 1

#### Sponsor

##### Name of organization / entity

University of social welfare and rehabilitation sciences

##### Full name of responsible person

Hamidreza Khorram Khorshid

##### Street address

Koodakyar, Daneshjoo Blv, Evin , Tehran

##### City

Tehran

##### Province

Tehran

##### Postal code

1985713834

##### Phone

+98 21 7173 2000

##### Email

webmaster@uswr.ac.ir

##### Web page address

#### Grant name

#### Grant code / Reference number

#### Is the source of funding the same sponsor organization/entity?

Yes

#### Title of funding source

University of social welfare and rehabilitation 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

University of social welfare and rehabilitation sciences

##### Full name of responsible person

Mehran Fathizadeh

##### Position

Researcher

##### Latest degree

Master

##### Other areas of specialty/work

Physiotherapy

##### Street address

Koodakyar Ave, Daneshjoo Blv, Evin, Tehran, Iran

##### City

Tehran

##### Province

Tehran

##### Postal code

0000000000

**Phone**  
+98 21 6609 8062  
**Fax**  
**Email**  
M3hran.fathi@gmail.com

## Person responsible for scientific inquiries

### Contact

**Name of organization / entity**  
University of social welfare and rehabilitation sciences  
**Full name of responsible person**  
Mehran Fathizadeh  
**Position**  
Researcher  
**Latest degree**  
Master  
**Other areas of specialty/work**  
Physiotherapy  
**Street address**  
Koodakyar Ave, Daneshjoo Blv, Evin, Tehran, Iran  
**City**  
Tehran  
**Province**  
Tehran  
**Postal code**  
0000000000  
**Phone**  
+98 21 6609 8062  
**Fax**  
**Email**  
M3hran.fathi@gmail.com

## Person responsible for updating data

### Contact

**Name of organization / entity**  
University of social welfare and rehabilitation sciences  
**Full name of responsible person**  
Mehran Fathizadeh  
**Position**  
Researcher  
**Latest degree**  
Master  
**Other areas of specialty/work**  
Physiotherapy  
**Street address**

Koodakyar Ave, Daneshjoo Blv, Evin, Tehran, Iran  
**City**  
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**Province**  
Tehran  
**Postal code**  
0000000000  
**Phone**  
+98 21 6609 8062  
**Fax**  
**Email**  
M3hran.fathi@gmail.com

## 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

No - There is not a plan to make this available

### Clinical Study Report

Yes - There is a plan to make this available

### Analytic Code

Not applicable

### 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

Collected clinical data

### When the data will become available and for how long

Immediately after article publishing

### To whom data/document is available

All interested people

### Under which criteria data/document could be used

To get more details of collected data

### From where data/document is obtainable

contact via : m3hran.fathi@gmail.com

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

NO special process. IT may take about 2 weeks from contacting via E-mail

### Comments