

Clinical Trial Protocol

Iranian Registry of Clinical Trials

28 Jun 2026

The effect of trigger point dry needling of Quadratus lumborum on improvement of maximal isometric force of gluteus medius , pain and function of patellofemoral joint in patients with patellofemoral pain syndrome (single-blind randomized controlled trial)

Protocol summary

Study aim

The effect of dry needling of quadratus lumbarum muscle technique on improvement of maximal isometric force of gluteus medius muscle, pain and patellofemoral joint function in patients with patellofemoral pain syndrome

Design

A randomized control trial with blinded examiner .52 patients , enrolled between February 2020 and June 2020.

Settings and conduct

This study done at Mahdiah Clinic and the participants randomly divided into two groups after filling out the consent form. Subjects in the intervention group receive 6 sessions of routine physiotherapy and 3 sessions of dry needle technique and in the control group receive only 6 sessions of routine physiotherapy. An evaluator who is blind to the grouping of patients and the treatment provided for each measure outcomes before the first session, immediately after the first session, and at the end of the sixth session.

Participants/Inclusion and exclusion criteria

Entry requirements: individuals between 18 and 40 years old; unilateral symptoms; gradual onset of pain not due to trauma and lasts for at least three months; anterior or knee pain that is aggravated by at least two of the following: _ Prolonged sitting _ Climbing stairs _ Squat _ Run _ to jump _ Lee Lee Exclusion criteria: a patella apprehension test or patella dislocation; knee effusion; radicular pain from hip or lumbar; surgery on the patellofemoral joint; signs of meniscus, collateral or cruciate ligament injury.

Intervention groups

The intervention group received 6 sessions of routine physiotherapy electrical stimulation and exercise therapy. In the first, third and fifth session, dry needle technique of the quadratus lumbarum is performed.

Control group will receive 6 sessions of routine physiotherapy including electrical stimulation and exercise therapy.

Main outcome variables

Maximal isometric force of gluteus medius; pain; function

General information

Reason for update

Acronym

IRCT registration information

IRCT registration number: **IRCT20200113046113N1**

Registration date: **2020-02-01, 1398/11/12**

Registration timing: **prospective**

Last update: **2020-02-01, 1398/11/12**

Update count: **0**

Registration date

2020-02-01, 1398/11/12

Registrant information

Name

Fatemeh Mehrabi

Name of organization / entity

Country

Iran (Islamic Republic of)

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Recruitment status

Recruitment complete

Funding source

Expected recruitment start date

2020-02-20, 1398/12/01

Expected recruitment end date

2020-06-19, 1399/03/30

Actual recruitment start date

empty

Actual recruitment end date

empty

Trial completion date

empty

Scientific title

The effect of trigger point dry needling of Quadratus lumborum on improvement of maximal isometric force of gluteus medius , pain and function of patellofemoral joint in patients with patellofemoral pain syndrome (single-blind randomized controlled trial)

Public title

The effect of Quadratus lumbarum muscle dry needling in patients with patellofemoral pain syndrome

Purpose

Treatment

Inclusion/Exclusion criteria**Inclusion criteria:**

Men and women Aged 18-40 years Unilateral Insidious onset of symptoms not due to trauma present for at least 3 months Anterior or Peripatellar knee pain during at least 2 of the following:Prolonged sitting,Stair ascent, squatting, running, jumping, hopscotch Ability to read and write The minimum visual scale being 2

Exclusion criteria:

Positive patellar apprehension sign and History of recurrent patellar dislocation Knee joint effusion Hip or lumbar referred pain Previous surgery to the patellofemoral joint Prolonged nonsteroidal anti-inflammatory drug or corticosteroid use Known articular cartilage damage based on XRay Local or systemic infections or inflammation Bleeding disorders Taking anti coagulation medication Needle phobia Pregnancy The history of regular exercise activity Leg length discrepancy Hip adductor shortness Neurological low back pain Meniscal, Cruciate or collateral ligament injury

Age

From **18 years** old to **40 years** old

Gender

Both

Phase

N/A

Groups that have been masked

- Outcome assessor

Sample size

Target sample size: **52**

Randomization (investigator's opinion)

Randomized

Randomization description

Subjects were randomly divided into control and experimental groups using randomized block design

Blinding (investigator's opinion)

Single blinded

Blinding description

The evaluator is not aware of what patients are in the group and what treatment is being given to them.

Placebo

Not used

Assignment

Parallel

Other design features**Secondary Ids**

empty

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

Ethics committee of Isfahan University of medical sciences

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Hezar Jerib street

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Isfahan

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Isfahan

Postal code

8174673461

Approval date

2020-01-02, 1398/10/12

Ethics committee reference number

IR.MUI.RESEARCH.REC.1398.570

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

Patellofemoral pain syndrome

ICD-10 code

M22.2X

ICD-10 code description

Patellofemoral disorders

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

Maximal isometric force of gluteus medius muscle

Timepoint

Before the first session, immediately after the first session, the end of the sixth session

Method of measurement

Handheld dynamometer

2**Description**

Pain

Timepoint

Before the first session, immediately after the first session, the end of the sixth session

Method of measurement

3

Description

Function of patellofemoral joint

Timepoint

Before the first session, immediately after the first session, the end of the sixth session

Method of measurement

Kujala scoring questionnaire

Secondary outcomes

empty

Intervention groups

1

Description

Intervention group: Six sessions routine physiotherapy include electrical stimulation performed by TENS and exercise therapy for knee joint . we also do Quadratus lumbarum muscle dry needling technique in first, third and fifth session.

Category

Rehabilitation

2

Description

Control group: Six sessions routine physiotherapy include electrical stimulation performed by TENS and exercise therapy for knee joint

Category

Rehabilitation

Recruitment centers

1

Recruitment center

Name of recruitment center

Mahdieh clinic

Full name of responsible person

Fatemeh Mehrabi

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Sponsors / Funding sources

1

Sponsor

Name of organization / entity

Esfahan University of Medical Sciences

Full name of responsible person

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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

Fatemeh Mehrabi

Position

Master student

Latest degree

Bachelor

Other areas of specialty/work

Physiotherapy

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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

Yes - There is a plan to make this available

Data Dictionary

Yes - There is a plan to make this available

Title and more details about the data/document

All obtained data, such as demographic information and primary outcomes measured at all time points, and data analysis can be shared after unidentifiable subjects.

When the data will become available and for how long

The access period begins after the results are published.

To whom data/document is available

Anyone working in the medical and rehabilitation field can access the information.

Under which criteria data/document could be used

After receiving information if they wish to use it in future studies, they should seek permission from the designers.

From where data/document is obtainable

Postal address: School of Rehabilitation Sciences, Hezar Jerib street, Isfahan, Iran Postcode: 8174673461 Email address : fatemehhh@rehab.mui.ac.ir fmehrabipt@yahoo.com Phone Number : +98 31 3792 3071 Cell Number : 09139649659 Responsible person: Fatemeh Mehrabi

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

The applicant can request the data by email. After 5 working days the data will be sent to them.

Comments