

Clinical Trial Protocol

Iranian Registry of Clinical Trials

09 Jun 2026

Comparison of the combined gluteus medius and quadratus lumborum dry needling & exercise with exercise alone on knee joint pain and function in female athletes with patellofemoral pain syndrome

Protocol summary

Study aim

The purpose of this study is to evaluate the effect of the effect of gluteus medius and quadratus lumborum muscle dry needling combined with therapeutic exercises on pain intensity, Pain threshold, function and balance in female athletes with patellofemoral pain syndrome.

Design

Two arm parallel group randomised trial with single-blind

Settings and conduct

In this clinical trial, individuals are included in the study if they are eligible and randomly assigned to one of the two intervention and control groups (each group of 20) using the Randomizer software

Participants/Inclusion and exclusion criteria

Women athletic patients with patellofemoral pain syndrome complaining gradual anterior knee pain during last 3 months and age between 18 to 45 years old are included. The patients with any previous knee trauma or other pathologies such as meniscus and ligament injuries are excluded.

Intervention groups

Minimal sample size for this study was determined as 20 patients in each group. Patients will randomly assigned either to control or intervention groups. Both groups will receive conventional physical therapy exercises for patellofemoral pain syndrome during 4 weeks (8 sessions), while the intervention group will receive gluteus medius and quadratus lumborum muscle dry needling.

Main outcome variables

Pain intensity

General information

Reason for update

Acronym

IRCT registration information

IRCT registration number: **IRCT20120411009440N20**

Registration date: **2018-08-02, 1397/05/11**

Registration timing: **prospective**

Last update: **2018-09-04, 1397/06/13**

Update count: **1**

Registration date

2018-08-02, 1397/05/11

Registrant information

Name

Soraya Pirouzi

Name of organization / entity

School of Rehabilitation Sciences, Shiraz University of Medical Sciences

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Recruitment status

Recruitment complete

Funding source

Expected recruitment start date

2018-08-05, 1397/05/14

Expected recruitment end date

2018-11-22, 1397/09/01

Actual recruitment start date

empty

Actual recruitment end date

empty

Trial completion date

empty

Scientific title

Comparison of the combined gluteus medius and quadratus lumborum dry needling & exercise with exercise alone on knee joint pain and function in female athletes with patellofemoral pain syndrome

Public title

The effect of gluteus medius and quadratus lumborum dry needling in patients with patellofemoral pain syndrome

Purpose

Treatment

Inclusion/Exclusion criteria**Inclusion criteria:**

Women aged 18 to 45 years old with unilateral or bilateral patellofemoral pain syndrome; anterior or retro patellar knee pain during at least 2 following activities: prolonged sitting, ascending or descending stair, squatting, kneeling, jumping and running; pain on palpation of medial and lateral patellar facets and positive patellar grind test; an insidious onset of symptoms not related to trauma for the last 3 months; Kujala scores less than 85 out of 100; numeric rate scale scores ranging 3 to 10 during last week. having trigger point in the gluteus medius muscle in the affected side and quadratus lumborum muscle in the non-affected side

Exclusion criteria:

Other knee joint pathologies such as meniscus and ligament injuries, osteoarthritis and tendon impairments; patellar subluxations or dislocations; Sinding-Larsen disease; Osgood-Schlatter disease; pelica syndrome; ankle and knee injury; hip joint pathologies such as ligament injuries, osteoarthritis and tendon impairments; subluxations or dislocations and hip injury; referral pain from other joints including lumbar spine, hip and sacroiliac joints and spinal fractures; LBP without source of myofascial; any pain during sitting such as Coccydynia; previous knee surgery; Structural and biomechanical problems such as: Varus and Valgus knee; overweight $32 < \text{BMI}$ Prohibition the use of dry needles such as: metabolic diseases such as diabetes, rheumatic diseases and neuromuscular diseases; pregnancy; Respiratory and peripheral vascular disease, cancer and any malignancy; Immune deficiency; Menstruation; needle phobia; Hemorrhagic diseases and the use of anticoagulant drugs; liver and kidney disease

Age

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

Gender

Female

Phase

N/A

Groups that have been masked

- Outcome assessor

Sample size

Target sample size: **40**

Randomization (investigator's opinion)

Randomized

Randomization description

Randomization tool: Statistical software

Blinding (investigator's opinion)

Single blinded

Blinding description

single blind, the assessor are not aware of the group

Placebo

Not used

Assignment

Parallel

Other design features**Secondary Ids**

empty

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

Ethics committee of Shiraz University of Medical Sciences

Street address

Zand Blvd., Central Building of Shiraz University of Medical Sciences

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Postal code

713451978

Approval date

2018-07-04, 1397/04/13

Ethics committee reference number

IR.SUMS.REC.1397.326

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

Patellofemoral disorders

ICD-10 code**ICD-10 code description****Primary outcomes****1****Description**

Pain threshold

Timepoint

first and 8th sessions and 2 weeks after the intervention

Method of measurement

algometry

2**Description**

Pain intensity

Timepoint

first and 8th sessions and 2 weeks after the intervention

Method of measurement

Numeric Rate Scale (NRS)

3

Description

Functional stepping

Timepoint

first and 8th sessions and 2 weeks after the intervention

Method of measurement

Step down test

4

Description

function

Timepoint

first and 8th sessions and 2 weeks after the intervention

Method of measurement

kujula questionnaire

5

Description

balance

Timepoint

first and 8th sessions and 2 weeks after the intervention

Method of measurement

Y-Balance Test

Secondary outcomes

empty

Intervention groups

1

Description

Intervention group: The same as control group plus gluteus medius and quadratus lumborum dry needling for 4 sessions, are given over a period of four weeks one session per week. The needle technique is fast in and out into the muscle tight band in three times

Category

Rehabilitation

2

Description

Control group: Exercise progressively, first week: Stretching exercises for quadriceps, hamstring (2set 15 repetition, 10 second hold); quadriceps setting in 10 degree terminal knee extension (3set 10 repetition), second week: sideling straight leg raises and hip abductors and external rotator strengthening exercise (Clamshells) (3 set, 15 repetition). third week: plank, side plank (3set 20 repetition). Fourth week: minisquat, minilung, single leg stance step down (Stairs with height 18 cm) (3set 25 repetition)

Category

Rehabilitation

Recruitment centers

1

Recruitment center**Name of recruitment center**

Physiotherapy clinic of Shiraz School of Rehabilitation

Full name of responsible person

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Sponsors / Funding sources

1

Sponsor**Name of organization / entity**

Shiraz 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

Shiraz 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

piroozis@sums.ac.ir

Contact

Name of organization / entity

Shiraz University of Medical Sciences

Full name of responsible person

Hanieh zarei

Position

Master science student of Physical Therapy

Latest degree

Bachelor

Other areas of specialty/work

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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

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

Clinical Study Report

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

Analytic Code

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

Data Dictionary

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