

# Clinical Trial Protocol

## Iranian Registry of Clinical Trials

10 Jun 2026

### Comparative effect of Functional Patellofemoral Joint Mobilizations and Open-Pack Patellofemoral Joint Mobilization for Patellofemoral Pain Syndrome; A Randomized Clinical Trial.

#### Protocol summary

##### Study aim

To compare functional patellofemoral joint mobilizations with the standard open-pack patellofemoral joint mobilization for reducing pain associated with patellofemoral pain syndrome.

##### Design

A single blinded quantitative study ,following the research design of randomized control trial. Randomization will be through coin tossing method. Active control group will be used to compare the results

##### Settings and conduct

Subjects with patellofemoral pain syndrome will be taken from these 3 hospital settings: Allied hospital Faisalabad, District Headquater hospital Faisalabad and Madinah Teaching Hospital Faisalabad. Blinding: participants will be blinded they will not be aware whether they are in treatment or control group as the participants of both the groups will receive 3 step intervention .Physiotherapist will provide different mobilization to each group which will be hidden from the subjects.

##### Participants/Inclusion and exclusion criteria

Inclusion criteria: Subjects having pain in the knee associated with, during or after the activity including stair ascending/descending, squatting, jumping, prolonged sitting etc. Non-inclusion criteria: Meniscal injury or any other articular injury; Knee ligament laxity; History of recent knee surgery; History of recent patellar dislocation.

##### Intervention groups

Intervention group 1: 1. Hot pack (10 min) . 2. Functional patellofemoral joint mobilizations(medial patellofemoral glide in 4 functional positions) 3. Baseline quadricep strengthening exercise program. Intervention group 2: 1.Hot pack (10 min) . 2.Baseline quadricep strengthening exercise program. 3.Grade I-11 Medial patellofemoral mobilizations (in one position only).

##### Main outcome variables

Visual Analogue Scale score; Anterior Knee Pain Scale score

#### General information

##### Reason for update

##### Acronym

##### IRCT registration information

IRCT registration number: **IRCT20220401054385N1**

Registration date: **2022-06-05, 1401/03/15**

Registration timing: **retrospective**

Last update: **2022-06-05, 1401/03/15**

Update count: **0**

##### Registration date

2022-06-05, 1401/03/15

##### Registrant information

##### Name

Laiba Nadeem

##### Name of organization / entity

The University of Faisalabad

##### Country

Pakistan

##### Phone

+92 308 6511400

##### Email address

dpt-fa14-097@tuf.edu.pk

##### Recruitment status

**Recruitment complete**

##### Funding source

##### Expected recruitment start date

2022-04-29, 1401/02/09

##### Expected recruitment end date

2022-05-30, 1401/03/09

##### Actual recruitment start date

empty

**Actual recruitment end date**

empty

**Trial completion date**

empty

**Scientific title**

Comparative effect of Functional Patellofemoral Joint Mobilizations and Open-Pack Patellofemoral Joint Mobilization for Patellofemoral Pain Syndrome; A Randomized Clinical Trial.

**Public title**

Effect of Patellofemoral Joint Mobilizations for knee pain

**Purpose**

Treatment

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

Patients at the age of 18-35 years old Both gender Willingness to participate in the research Willingness for random allocation to either of the two groups Pain in the knee associated with, during or after the activity including stair ascending/descending, squatting, jumping, prolonged sitting etc Insidious onset of pain that is unrelated to trauma Patient having at least one positive test from these physical tests: 1)Patellofemoral grind test. 2) Patellar apprehension test.3) Step down test Visual analogue scale rating of anterior knee pain during daily activities at a minimum of 30 mm to 70 mm on 100 mm scale

**Exclusion criteria:**

Meniscal injury or any other articular injury. Knee ligament laxity. History of recent knee surgery History of recent patellar dislocation NSAID or corticosteroid drug use before testing within 24 hours

**Age**

From **18 years** old to **35 years** old

**Gender**

Both

**Phase**

N/A

**Groups that have been masked**

- Participant

**Sample size**

Target sample size: **50**

**Randomization (investigator's opinion)**

Randomized

**Randomization description**

Sample will be allocated to Group A (Functional mobilization) and Group B (Open pack mobilization) randomly using coin tossing. Method of randomization: Simple randomization Tool for randomization: Sealed envelopes Unit of randomization: Individual Allocation will be concealed from the participants, they will not know whether they are in the treatment or control group,

**Blinding (investigator's opinion)**

Single blinded

**Blinding description**

Participants of both groups will be unaware whether they are in active control group or the treatment group after taking informed consent from them. Participants of both groups will receive the same baseline treatment that is

quadriceps strengthening exercises. After that Physiotherapist will apply different types of knee mobilizations to both groups of which the participants will be unaware of.

**Placebo**

Not used

**Assignment**

Other

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

empty

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

Ethics committee of The University of Faisalabad

**Street address**

4-km,Sargodha Road, University Town, Faisalabad, Pakistan.

**City**

Faisalabad

**Postal code**

38000

**Approval date**

2022-04-07, 1401/01/18

**Ethics committee reference number**

TUF/DR/MSPP/322

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

Patellofemoral pain syndrome

**ICD-10 code**

M22.2

**ICD-10 code description**

Patellofemoral disorders

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

Pain in the anterior knee

**Timepoint**

Pain level will be measured before applying the intervention and then after 4 weeks after the completion of intervention.

**Method of measurement**

Visual Analogue scale and anterior knee pain scale will be used to measure the pain.

**Secondary outcomes**

empty

## Intervention groups

### 1

#### Description

Intervention group (Group A) ; 1. Hot pack (10 min)  
2. Grade 1-11 Functional patellofemoral joint mobilizations  
3. Baseline Quadriceps strengthening exercise program.

#### Category

Rehabilitation

### 2

#### Description

Control group: Hot pack (10 min)  
2. Grade 1-11 open-pack patellofemoral joint mobilizations  
3. Baseline Quadriceps strengthening exercise program.

#### Category

Rehabilitation

## Recruitment centers

### 1

#### Recruitment center

##### Name of recruitment center

Madinah Teaching Hospital

##### Full name of responsible person

Dr. Sidra Majeed

##### Street address

Sargodha Rd, University Town, Faisalabad, Punjab,  
Pakistan.

##### City

Faisalabad

##### Postal code

38000

##### Phone

+92 334 7731241

##### Email

info@mth.org.pk

### 2

#### Recruitment center

##### Name of recruitment center

Allied Hospital Faisalabad

##### Full name of responsible person

Dr. Sidra Majeed

##### Street address

Jail Road adjacent Faisalabad Medical University, Near  
Sargodha Road, Faisalabad, Pakistan

##### City

Faisalabad

##### Postal code

38000

##### Phone

+92 334 7731241

##### Email

sidra.majeed@tuf.edu.pk

### 3

#### Recruitment center

##### Name of recruitment center

DHQ hospital Faisalabad

##### Full name of responsible person

Dr. Sidra Majeed

##### Street address

Near Railway Station, Mall Road, Faisalabad, Pakistan

##### City

Faisalabad

##### Postal code

38000

##### Phone

+92 334 7731241

##### Email

sidra.majeed@tuf.edu.pk

## Sponsors / Funding sources

### 1

#### Sponsor

##### Name of organization / entity

The University of Faisalabad

##### Full name of responsible person

Dr. Sidra Majeed, PT

##### Street address

4-km Sargodha road, University Town, Faisalabad,  
Pakistan

##### City

Faisalabad

##### Postal code

38000

##### Email

sidra.majeed@tuf.edu.pk

#### Grant name

#### Grant code / Reference number

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

No

#### Title of funding source

The University of Faisalabad

#### Proportion provided by this source

100

#### Public or private sector

Private

#### Domestic or foreign origin

Domestic

#### Category of foreign source of funding

*empty*

#### Country of origin

#### Type of organization providing the funding

Other

## Person responsible for general inquiries

#### Contact

##### Name of organization / entity

The University of Faisalabad

##### Full name of responsible person

Dr. Sidra Majeed, PT

**Position**

Associate Professor

**Latest degree**

Master

**Other areas of specialty/work**

Physiotherapy

**Street address**

4-km Sargodha road, University town, Faisalabad,  
Pakistan

**City**

Faisalabad

**Province**

Punjab

**Postal code**

38000

**Phone**

+92 334 7731241

**Email**

sidra.majeed@tuf.edu.pk

**Latest degree**

Master

**Other areas of specialty/work**

Physiotherapy

**Street address**

House no 12,sahil homes, mian zulfiqar ali shahid  
road, Faisalabad

**City**

Faisalabad

**Province**

Punjab

**Postal code**

38000

**Phone**

+92 308 6511400

**Fax****Email**

dpt-fa14-097@tuf.edu.pk

**Person responsible for scientific inquiries****Contact****Name of organization / entity**

The University of Faisalabad

**Full name of responsible person**

Laiba Nadeem

**Position**

Student

**Latest degree**

Master

**Other areas of specialty/work**

Physiotherapy

**Street address**

House no 12,sahil homes, mian zulfiqar ali shahid  
road, Faisalabad

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**Postal code**

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**Phone**

+92 308 6511400

**Fax****Email**

dpt-fa14-097@tuf.edu.pk

**Person responsible for updating data****Contact****Name of organization / entity**

The University of Faisalabad

**Full name of responsible person**

Laiba Nadeem

**Position**

Student

**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**

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

**Title and more details about the data/document**

All collected deidentified participant data set for the  
outcome measures.

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

4 months after publication

**To whom data/document is available**

People working in the academic institutions

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

IPD and any any additional supporting documents will be  
provided for the research on the similar topic and can be  
obtained by asking through the contact person  
mentioned below

**From where data/document is obtainable**

Contact person: Laiba Nadeem contact no: 0308  
6511400 email:laiba\_nadeem1708@gmail.com

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

Data files will be provided after talking to the contact  
person as mention in the above slot.

**Comments**