Treatment of knee pain with combined instrument-assisted soft tissue mobilization

Submission date 28/01/2023	Recruitment status No longer recruiting	 Prospectively registered Protocol
Registration date 07/02/2023	Overall study status Completed	 Statistical analysis plan [X] Results
Last Edited 02/05/2025	Condition category Musculoskeletal Diseases	Individual participant data

Plain English summary of protocol

Background and study aims

This study looks at the effects of instrument-assisted soft tissue mobilization (IASTM) therapy on patellofemoral pain syndrome, a common knee condition which causes pain at the front of the knee around the area known as the patellofemoral joint. The study aims to find out if the therapy reduces patellofemoral discomfort, and improves function, strength, and joint mobility.

Who can participate? Adult patients aged between 18 and 35 years old with patellofemoral discomfort

What does the study involve?

This study lasted for four weeks and was conducted at the Experimental Centre for Sports Rehabilitation of the Wuhan Sports University. To help the trial run smoothly, the study also included resistance training or the use of tui na, a traditional Chinese massage technique.

What are the possible benefits and risks of participating?

The ability to help people with patellofemoral discomfort, reduce or eliminate knee pain, boost lower limb strength, and regain lower limb function is the potential benefit of the project. While there is a danger that the procedure will cause soft tissue damage, this risk is low and the experiment is entirely under control.

Where is the study run from? Wuhan Sports University (China)

When is the study starting and how long is it expected to run for? May 2022 to October 2022

Who is funding the study? Wuhan Sports University (China)

Who is the main contact? Prof Lianqing Wu, 784389072@qq.com

Contact information

Type(s) Public

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Additional identifiers

EudraCT/CTIS number Nil known

IRAS number

ClinicalTrials.gov number Nil known

Secondary identifying numbers Nil known

Study information

Scientific Title

Treatment of patellofemoral joint pain with combined instrument-assisted soft tissue mobilization

Study objectives

Instrument-assisted soft tissue mobilization treatment can improve the symptoms of patients with patellofemoral joint pain, including pain, function and lower limb strength.

Ethics approval required

Ethics approval required

Ethics approval(s)

1. Approved 02/06/2022, Medical Ethics Committee of Wuhan Institute of Sports (No. 461 Luoyu Road, Hongshan District, Wuhan, 430079, China; +86 18935896605; liuyang121795@qq.com), ref: 2022062

2. Submitted 01/09/2022, Medical Human Ethics Review Committee of Wuhan Institute of Physical Education (Wuhan Sports University, Wuhan, 430000, China; +86 18935836605; liuyang121795@qq.com), ref: N202061

Study design Randomized controlled study

Primary study design Interventional

Secondary study design Randomised controlled trial

Study setting(s) School

Study type(s) Treatment

Participant information sheet

Not available in web format, please use the contact details to request a participant information sheet

Health condition(s) or problem(s) studied

Patellofemoral joint pain

Interventions

Patients with patellofemoral joint pain were randomly divided into an instrument-assisted soft tissue mobilization (IASTM) test group and a control group. The IASTM test group was treated with IASTM combined with resistance training while the control group received only IASTM. Both groups were treated for 4 weeks.

At four time points, before treatment, after the first treatment, after the second treatment and after the third treatment, the visual analog pain score (VAS), the Lysholm score of the knee and the modified Thomas test (MTT) were compared and analyzed.

Before and 4 weeks after treatment, the maximum isometric muscle strength test system of the lower limb extensor was undertaken.

Intervention Type

Procedure/Surgery

Primary outcome measure

1. Pain measured using the visual analogue pain score (VAS) assessed before the start of the experiment, and in the first and last weeks of the experiment

2. Knee-specific symptoms measured using the Lysholm Knee Scoring System assessed before the start of the experiment, and in the first and last weeks of the experiment

3. Flexibility of the iliopsoas, rectus femoris and tensor fascia latae (hip flexion contracture and hip extensibility) measured using the modified Thomas test (MTT) assessed before the start of the experiment, and in the first and last weeks of the experiment

4. Muscle strength test measured using the maximum isometric muscle strength test system of

the lower limb extensor assessed before the start of the experiment and in the last week of the experiment

Secondary outcome measures There are no secondary outcome measures

Overall study start date 11/05/2022

Completion date 19/10/2022

Eligibility

Key inclusion criteria

Aged between 18 and 35 years old
 Be able to cooperate with rehabilitation physiotherapy, training and follow-up, and will not withdraw from the study without reason

Participant type(s)

Patient

Age group Adult

Lower age limit 18 Years

Sex Both

Target number of participants 46

Total final enrolment 38

Key exclusion criteria

1. Exogenous injury of knee joint, including bruise, hit, cut, scald, contusion, etc., without external or subcutaneous bleeding due to skin rupture

2. Abnormal natural bone structure

3. Inflammation, patellar dislocation or subluxation, knee ligament injury

4. History of arthroscopic surgery of knee joint within one year

5. Injury or discomfort of other body parts

Date of first enrolment

01/09/2022

Date of final enrolment 18/09/2022

Locations

Countries of recruitment China

Study participating centre Wuhan Sports University No. 461 Luoyu Road Hubei Province Wuhan City China 430079

Sponsor information

Organisation Wuhan Sports University

Sponsor details Wuhan Institute of Physical Education Wuhan China 430000 +861 027-87191823 Webmaster@whsu.edu.cn

Sponsor type University/education

Website http://lxwten.whsu.edu.cn/

ROR https://ror.org/004je0088

Funder(s)

Funder type University/education

Funder Name

Results and Publications

Publication and dissemination plan

Planned publication in a high-impact peer-reviewed journal

Intention to publish date

06/07/2023

Individual participant data (IPD) sharing plan

The data set generated during this study is not expected to be available because these data relate to the personal privacy of the subjects.

IPD sharing plan summary

Not expected to be made available

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient- facing?
<u>Results</u> article	Ccomparative study of the efficacy of instrument-assisted soft tissue mobilization and massage techniques	13/11 /2023	01/12 /2023	Yes	No
<u>Results</u> article	Effect of combined IASTM and blood flow treatment	31/08 /2023	02/05 /2025	Yes	No